

A faunal inventory of methane seeps on the Pacific margin of Costa Rica

Charlotte A. Seid¹, Avery S. Hiley¹, Marina F. McCowin¹, José I. Carvajal¹, Harim Cha¹,
Shane T. Ah Yong^{2,3}, Oliver S. Ashford⁴, Odalysa Breedy⁵, Douglas J. Eernisse⁶, Shana K. Goffredi⁷,
Michel E. Hendrickx⁸, Kevin M. Kocot⁹, Christopher L. Mah¹⁰, Allison K. Miller¹¹,
Nicolás Mongiardino Koch¹, Rich Mooi¹², Timothy D. O'Hara¹³, Fredrik Pleijel¹⁴, Josefin Stiller¹⁵,
Ekin Tilic¹⁶, Paul Valentich-Scott¹⁷, Anders Warén¹⁸, Mary K. Wicksten¹⁹, Nerida G. Wilson^{1,20,21},
Erik E. Cordes²², Lisa A. Levin¹, Jorge Cortés⁵, Greg W. Rouse¹

- 1 Scripps Institution of Oceanography, University of California San Diego, La Jolla, California, USA
 - 2 Australian Museum, Sydney, New South Wales, Australia
 - 3 University of New South Wales, Kensington, New South Wales, Australia
 - 4 Ocean Program, World Resources Institute, London, UK
 - 5 Universidad de Costa Rica, San José, Costa Rica
 - 6 California State University Fullerton, Fullerton, California, USA
 - 7 Occidental College, Los Angeles, California, USA
 - 8 Instituto de Ciencias del Mar y Limnología, Universidad Nacional Autónoma de México, Mazatlán, Sinaloa, Mexico
 - 9 University of Alabama, Tuscaloosa, Alabama, USA
 - 10 Smithsonian National Museum of Natural History, Washington, DC, USA
 - 11 University of Otago, Dunedin, New Zealand
 - 12 California Academy of Sciences, San Francisco, California, USA
 - 13 Museums Victoria, Melbourne, Victoria, Australia
 - 14 University of Gothenburg, Gothenburg, Sweden
 - 15 University of Copenhagen, Copenhagen, Denmark
 - 16 Senckenberg Research Institute and Natural History Museum, Frankfurt, Germany
 - 17 Santa Barbara Museum of Natural History, Santa Barbara, California, USA
 - 18 Swedish Museum of Natural History, Stockholm, Sweden
 - 19 Texas A&M University, College Station, Texas, USA
 - 20 Collections & Research, Western Australian Museum, Welshpool, Western Australia, Australia
 - 21 School of Biological Sciences, University of Western Australia, Perth, Western Australia, Australia
 - 22 Temple University, Philadelphia, Pennsylvania, USA
- Corresponding authors: Charlotte A. Seid (cseid@ucsd.edu); Greg W. Rouse (grouse@ucsd.edu)



Academic editor: Andrew Davinack
Received: 14 August 2024
Accepted: 7 October 2024
Published: 3 January 2025

ZooBank: <https://zoobank.org/ED8980D6-922C-47CF-AF1D-456B2EDC133A>

Copyright: © Charlotte A. Seid et al.
This is an open access article distributed under
terms of the Creative Commons Attribution
License ([Attribution 4.0 International – CC BY 4.0](https://creativecommons.org/licenses/by/4.0/)).

Abstract

The methane seeps on the Pacific margin of Costa Rica support extensive animal diversity and offer insights into deep-sea biogeography. During five expeditions between 2009 and 2019, we conducted intensive faunal sampling via 63 submersible dives to 11 localities at depths of 300–3600 m. Based on these expeditions and published literature, we compiled voucher specimens, images, and 274 newly published DNA sequences to present a taxonomic inventory of macrofaunal and megafaunal diversity with a focus on invertebrates. In total 488 morphospecies were identified, representing the highest number of distinct morphospecies published from a single seep or vent region to date. Of these, 131 are described species, at least 58 are undescribed species, and the remainder include some degree of taxonomic uncertainty, likely representing additional undescribed species. Of the described species, 38 are known only from the Costa Rica seeps and their vicinity. Fifteen range extensions are also reported for species known from Mexico, the Galápagos

Citation: Seid CA, Hiley AS, McCowin MF, Carvajal JI, Cha H, Ah Yong ST, Ashford OS, Breedy O, Eernisse DJ, Goffredi SK, Hendrickx ME, Kocot KM, Mah CL, Miller AK, Mongiardino Koch N, Mooi R, O'Hara TD, Pleijel F, Stiller J, Tilic E, Valentich-Scott P, Warén A, Wicksten MK, Wilson NG, Cordes EE, Levin LA, Cortés J, Rouse GW (2025) A faunal inventory of methane seeps on the Pacific margin of Costa Rica. ZooKeys 1222: 1–250. <https://doi.org/10.3897/zookeys.1222.134385>

seamounts, Chile, and the western Pacific; as well as 16 new depth records and three new seep records for species known to occur at vents or organic falls. No single evolutionary narrative explains the patterns of biodiversity at these seeps, as even morphologically indistinguishable species can show different biogeographic affinities, biogeographic ranges, or depth ranges. The value of careful molecular taxonomy and comprehensive specimen-based regional inventories is emphasized for biodiversity research and monitoring.

Key words: Biodiversity, biogeography, Central America, chemosynthetic ecosystem, COI, deep sea, DNA 'barcodes', molecular taxonomy, review

Table of contents

Introduction	3
Materials and methods.....	7
Sampling locations.....	7
Specimen collection and vouchering	7
Taxonomic scope	8
Specimen identification.....	10
DNA extraction and sequencing	10
Haplotype networks.....	12
Scanning electron microscopy (SEM)	12
Permits	12
Taxonomic listing	12
Annelida	13
Nemertea	66
Brachiopoda.....	69
Mollusca	69
Bryozoa	115
Entoprocta	115
Platyhelminthes.....	115
Chaetognatha	117
Nematoda	117
Arthropoda.....	117
Chordata	154
Echinodermata	156
Hemichordata	181
Xenacoelomorpha	182
Cnidaria	182
Porifera.....	200
Chromista	201
Ciliophora	202
Results	202
Taxonomic identification.....	202
New records	203
Biogeographic ranges	203
Occurrences at other chemosynthesis-based habitats	203
Discussion	206
Biodiversity at the CRM seeps	206

Comparison to checklists for Costa Rica.....	206
Comparison to global seep checklists	207
Comparison to seep-adjacent background habitats	207
Comparison to other seep, vent, and organic fall sites	207
Depth ranges at the CRM seeps	208
Range extensions reinforce eastern Pacific biogeographic connections ...	208
Other biogeographic connections	210
Intersection of seeps, vents, and organic remains.....	211
Overlap of species occurrences	211
Modes of symbiosis	212
Areas for further work	212
Conclusions.....	213
Importance of regional taxonomic inventories.....	213
Recommendations	214
Acknowledgements	214
Additional information.....	215
References.....	216
Supplementary material 1	250

Introduction

The Costa Rica margin (CRM) occupies a central position in the biogeographic and tectonic landscape of the eastern Pacific. The subduction of the Cocos Plate beneath the Caribbean Plate at the Middle America Trench gives rise to deep-sea methane seeps (hereafter called “seeps”) (Kahn et al. 1996; McAdoo et al. 1996; Zuleger et al. 1996), as are found on the neighboring continental margins from Oregon to Chile (Sibuet and Olu 1998; Van Dover et al. 2002; Levin et al. 2016). The boundaries of the Cocos Plate adjoin the hydrothermal vent fields of the East Pacific Rise and the Galápagos Rift, which also directly links to the CRM via the Coco Submarine Volcanic Range (Kimura et al. 1997; von Huene et al. 2000; Fiedler and Lavín 2006; Beaulieu et al. 2013). This geographic position suggests the potential for population connectivity to seeps north and south along the continental margins, to the tectonically adjacent vent fields, and to the central and western Pacific via deep-ocean equatorial circulation (Tunnicliffe et al. 1998; Van Dover et al. 2002; Bachraty et al. 2009; Miloslavich et al. 2011). Furthermore, the biogeographic history of the CRM has been influenced by the emergence of the Isthmus of Panama (Central American Isthmus) (Cortés and Wehrmann 2009), from the closure of deep-water exchange to the western Atlantic approximately 9 million years ago to the formation of the land barrier ~ 2.8 million years ago (O’Dea et al. 2016). The faunal diversity of the CRM thus offers a range of insights into deep-sea biogeography and evolution.

Active seepage of methane-rich fluids occurs at more than 100 sites located ca 50 km offshore along the CRM, from southern Nicaragua to the Osa Peninsula (Kahn et al. 1996; McAdoo et al. 1996; Zuleger et al. 1996; Kimura et al. 1997; Bohrmann et al. 2002; Sahling et al. 2008). The seeps occur at depths of ~ 400–3800 m at diverse geological features including mounds, faults, seamount subduction scars, and landslides (Kahn et al. 1996; McAdoo et al. 1996; Bohrmann et al. 2002; Sahling et al. 2008). The sites studied in this work are

shown in Fig. 1. The seeps also span hydrographic gradients. Between 400 and 1800 m, the temperature ranges from 9.5 to 2.7 °C and oxygen concentration ranges from 0.04 to 1.9 ml/l, with an oxygen minimum zone at 300–600 m (Bohrmann et al. 2002; Levin et al. 2015). Methane concentrations vary with depth and among sites across three orders of magnitude, from background levels of ~ 2 nmol/l to maximum levels of 75–1506 nmol/l (Bohrmann et al. 2002; Mau et al. 2014). These abiotic gradients invite careful comparisons of biodiversity across the ensemble of seep sites.

The first submersible dives at the CRM in 1994 revealed biological indicators of chemosynthetic activity, namely the presence of authigenic carbonates, microbial mats, and symbiont-bearing invertebrate megafauna (Fig. 2A–E) (Kahn et al. 1996; McAdoo et al. 1996). Authigenic carbonates are precipitated via anaerobic oxidation of methane (Ritger et al. 1987), in which a consortium of methane-oxidizing (methanotrophic) archaea and sulfate-reducing bacteria generates bicarbonate and hydrogen sulfide (Boetius et al. 2000; Orphan et al. 2001). Hydrogen sulfide, in turn, supports chemosynthesis by sulfur-oxidizing (thiotrophic) microbes (Tunnicliffe et al. 2003). At the CRM seeps, thiotrophic bacteria constitute many of the microbial mats (Mau et al. 2006, 2014; Bailey et al. 2011; Bernardino et al. 2012; Niemann et al. 2013) and play an important role as nutritional symbionts of vestimentiferan tubeworms, bathymodiolin mussels, and vesicomyid clams (Mau et al. 2006; Brzechffa and Goffredi 2021). As at other seeps (Suess et al. 1985; Sibuet and Olu 1998), these large-bodied symbiont-bearing invertebrate groups tend to dominate the biomass, and much of the initial biological characterization of the CRM seeps focused on these conspicuous megafauna (Kahn et al. 1996; Peek et al. 2000; Bohrmann et al. 2002; Goffredi et al. 2003).

The CRM seeps also harbor a variety of animals that do not directly depend on chemosynthetic symbionts for nutrition. Initial records of such non-obligate seep fauna included limpets, snails, crabs, galatheoids, crinoids, actinarians, corals, ophiuroids, echinoids, holothuroids, sponges, and macrurid fish (Kahn et al. 1996; Bohrmann et al. 2002). Further sampling has revealed diverse macrofaunal assemblages associated with carbonates and sediments at different habitat types (e.g., Levin et al. 2012, 2015; Ashford et al. 2021b; Pereira et al. 2021). To date, 48 animal species have been described from the CRM seeps, including annelids (Aguado and Rouse 2011; Borda et al. 2013; Summers et al. 2014b; Watson et al. 2016; McCowin and Rouse 2018; Rouse et al. 2018; Lindgren et al. 2019; Hatch et al. 2020; Salazar-Vallejo 2020a; Yen and Rouse 2020; Rouse and Kupriyanova 2021; Pearson and Rouse 2022; Vilalobos-Guerrero et al. 2024), corals (Opresko and Breedy 2010; Breedy et al. 2019), crustaceans (Thurber et al. 2011; Martin et al. 2018; Rodríguez-Flores et al. 2023), fish (Frale et al. 2023), echinoderms (Payne et al. 2023), mollusks (Barry and Kochevar 1999; Krylova and Sahling 2006; Martin and Goffredi 2012; Warén and Rouse 2016; McCowin et al. 2020), and nemerteans (Sagorny et al. 2022), as well as a ciliate (Pasulka et al. 2017) for a total of 49 eukaryotic species.

Furthermore, the CRM seeps show environmental and biological connections to other chemosynthesis-based habitats. The Jacó Scar “hydrothermal seep” site at 1800 m depth (Fig. 2F) appears to represent an intermediate

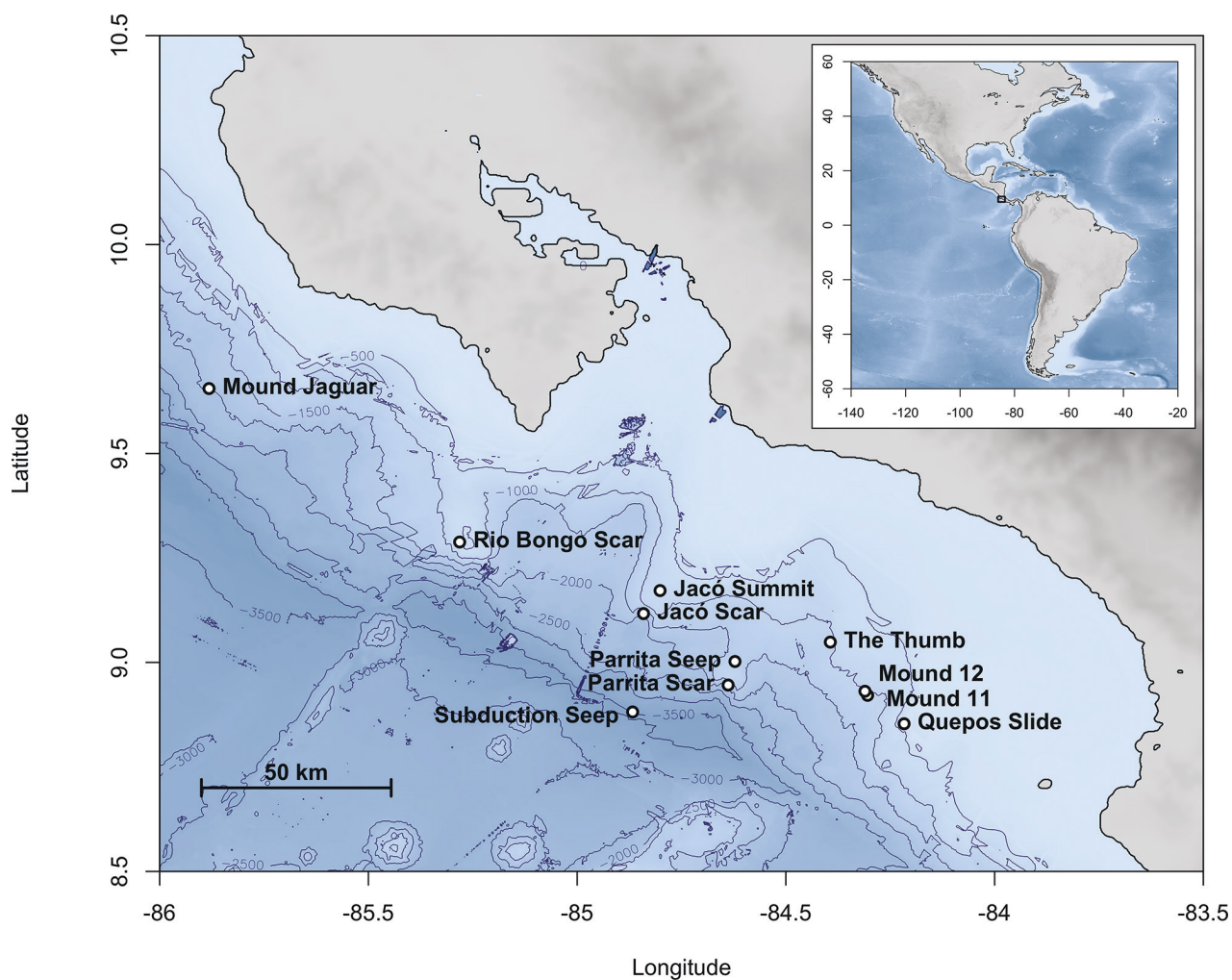


Figure 1. Map of the Costa Rica seep sites sampled in this study. Maps were generated using the R package *map* (Pante and Simon-Bouhet 2013) and bathymetric data from NOAA (<https://www.ncei.noaa.gov/maps/autogrid/>, accessed 25 August 2021).

environment between seeps and hydrothermal vents, as elevated water temperatures (up to 5.2 °C, representing nearly 3 °C above ambient) support certain vent-affiliated fauna (Levin et al. 2012). Wood falls (Fig. 2G) and animal carcasses (Fig. 2H) occur in the vicinity of these seeps, as might be expected from their proximity to a forested coast and its river discharges. Experimental deployments of wood and bone substrates have enabled further investigation of seep-adjacent organic fall communities (Hatch et al. 2020; Pereira et al. 2022; Payne et al. 2023). These intersectional habitats offer the opportunity to address questions of colonization, habitat suitability, and biogeography across different chemosynthesis-based ecosystems.

During five research cruises to the CRM in 2009–2019, we used deep submergence vehicles to conduct intensive faunal sampling. Based on these expeditions and published literature, we compiled voucher specimen records, images, and DNA sequences to present a taxonomic inventory of faunal diversity at these seeps. We report new range extensions and depth records, discuss connections to other chemosynthesis-based ecosystems, and assess biogeographic patterns.

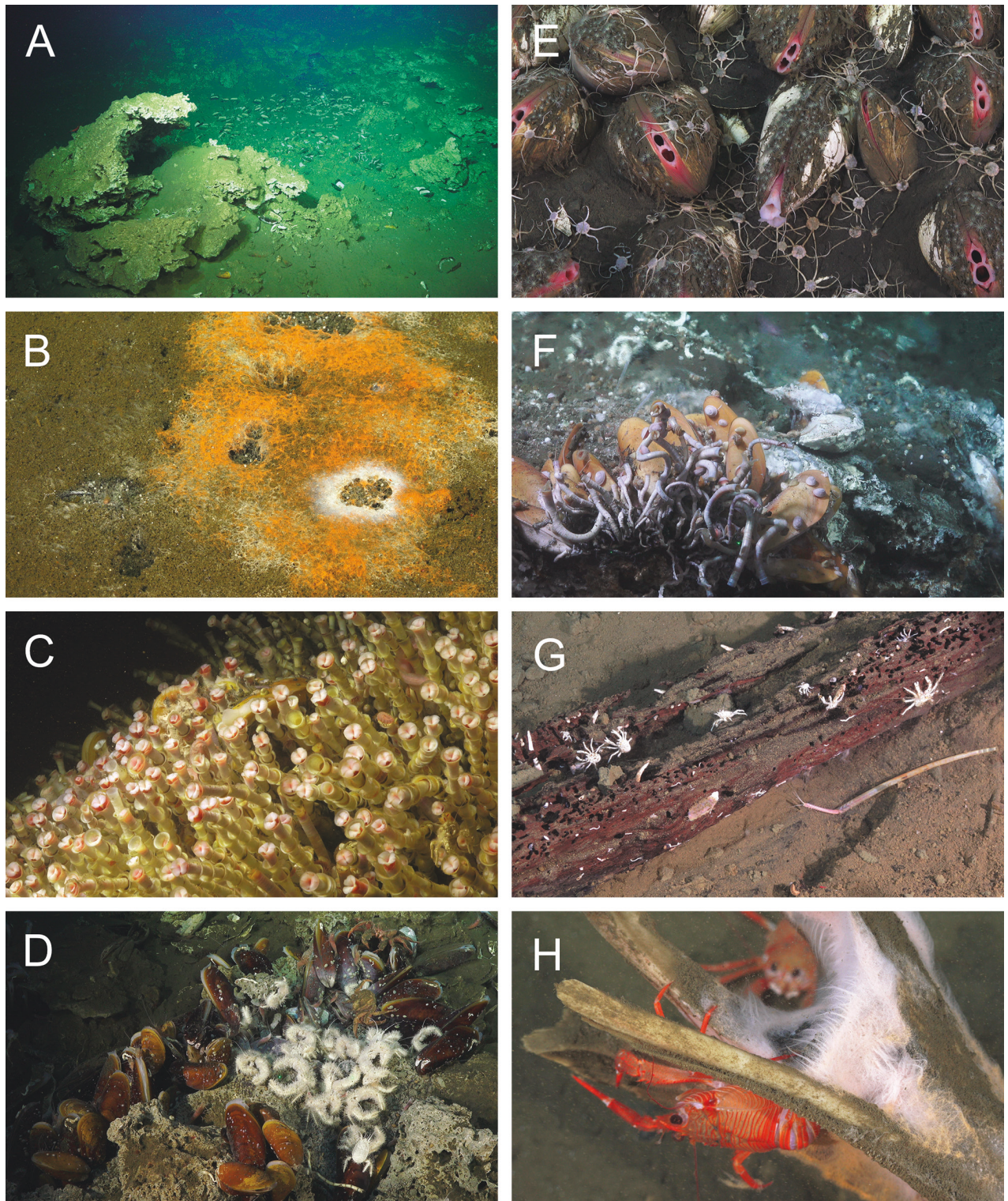


Figure 2. Diversity of habitats at the Costa Rica seeps. Credit: ROV SuBastian/Schmidt Ocean Institute **A** authigenic carbonates at Mound 12 (1006 m, Dive S0215) **B** microbial mat at Rio Bongo Scar (609 m, Dive S0219) **C** vestimentiferan tubeworm aggregation (predominantly *Lamellibrachia barhami*) at Jacó Scar (1814 m, Dive S0212) **D** mussels (*Bathymodiolus nancyschneiderae*) and yeti crabs (*Kiwa puravida*) on authigenic carbonates at Mound 12 (1006 m, Dive S0215) **E** vesicomyid clams at Jacó Scar (1781 m, Dive S0214) **F** seepage of higher-temperature fluid at the Jacó Scar hydrothermal seep site (1803 m, Dive S0214) **G** wood fall at Jacó Scar (1875 m, Dive S0214) **H** animal fall (billfish skull, utilized by *Grimothea monodon* squat lobsters) at Quepos Slide (403 m, Dive S0216).

Materials and methods

Sampling locations

Specimens were collected during five research cruises to the central CRM (Fig. 1): R/V Atlantis with DSV Alvin AT15-44 (2009), AT15-59 (2010), AT37-13 (2017), AT42-03 (2018); R/V Falkor with ROV SuBastian FK190106 (2019). Collection localities, depths, dates, and details are summarized in Table 1.

The submersible dives in this study primarily investigated areas of active methane seepage, as indicated by the presence of non-sedimented authigenic carbonates, microbial mats, or symbiont-bearing megafauna (Levin et al. 2015; Pasulka et al. 2017; Ashford et al. 2021b; Pereira et al. 2021). Certain dives also covered adjacent deep-sea habitat, as defined by the absence of these indicators, and “transition” habitat, characterized by partially sedimented authigenic carbonates, the remains of symbiont-bearing bivalves, and reduced density of seep-associated megafauna (Ashford et al. 2021b; Pereira et al. 2021). The complex biological and geochemical influences of seeps are thought to extend for hundreds of meters up to ~ 1 km in three dimensions (Levin et al. 2016; Ashford et al. 2021b), a spatial extent comparable to that covered by a given submersible dive in this study (typically a maximum radius of 1 km, centered at an active seep site). In a recent ecological study of the transition zone between seeps and adjacent deep-sea habitat at the CRM (Ashford et al. 2021b), the distances between accepted “active” and “transition” or “background” habitats ranged from 257–967 m. Therefore, all specimens collected on seep-focused dives and within ~ 500 m of active seepage were considered within the scope of this study, including a few opportunistically collected specimens from habitats that likely represent background or transition zones (e.g., three multicore deployments in Table 1). Non-seep habitats such as seamounts and Quepos Plateau (von Huene et al. 2000) were investigated during some of the same cruises but are outside the scope of this work.

We generally follow the locality names listed in Sahling et al. (2008). The locality named here as Parrita Seep has been variously cited as “Mound Quepos,” “Quepos Seep,” or “Quepos Seep at Parrita Scar” in previous work (e.g., Han et al. 2004; Sahling et al. 2008; Levin et al. 2015; McCowin and Rouse 2018). Based on the geological categorization of this site as a fault-controlled seep rather than a mound (Sahling et al. 2008) and its proximity to a separate site known as Parrita Scar, we follow other studies (McCowin et al. 2020; Rojas-Jimenez et al. 2020; Ashford et al. 2021b; Rouse and Kupriyanova 2021) in synonymizing these names as Parrita Seep.

Specimen collection and vouchering

Most specimens were collected during submersible dives using equipment such as hydraulic arms, suction samplers, scoops, push cores, a Bushmaster Jr. device for sampling tubeworm aggregations (Bergquist et al. 2002), and a customized mussel pot device for sampling mussel aggregations (Cordes et al. 2010). Some specimens were collected opportunistically during multicore deployments or plankton tows adjacent to the main study sites. Some specimens were collected from experimentally deployed substrates such as wood,

bone, or carbonate rocks, including rocks experimentally transplanted across different zones at the same locality and similar depth (Pereira et al. 2022), but no animals were transplanted between localities. Associations with wood or bone are indicated in each taxonomic listing, and specimens were collected from naturally occurring seep substrates unless otherwise indicated.

Specimens were maintained alive in chilled seawater, treated with an appropriate relaxing agent, and processed following recommended practices for DNA taxonomy of marine invertebrates (Gemeinholzer et al. 2010; Templado et al. 2010; Glover et al. 2015; Rouse et al. 2022). Live specimens were photographed using a handheld camera (e.g., Canon EOS M5, Canon Rebel T1, Nikon D70, Panasonic DMC-TS4, Ricoh WG-4, Ricoh WG-50) or a photomicroscopy station (e.g., a Leica S8 Apo or MZ9.5 stereomicroscope with a camera attachment for a Canon EOS Rebel T3i, T6s, or T6i; EOS M5; or PowerShot G9). Tissue subsamples or small whole specimens were preserved in 95% ethanol for genetic analysis. Voucher specimens for morphological analysis were typically fixed in 10% seawater formalin for at least 24 h, rinsed with fresh water, and transferred to 50% ethanol for long-term archival. For certain taxa, e.g., echinoderms and crustaceans, large voucher specimens were instead treated with 95% ethanol in several washes for at least 24 h each and then transferred to 50% ethanol for long-term archival. Selected specimens or tissues were treated with RNAlater (Ambion, Austin, TX), paraformaldehyde, glutaraldehyde, or osmium tetroxide. Some specimens from the 2009 and 2010 cruises were treated with the glyoxal fixative Prefer (Anatech Ltd., Battle Creek, MI) for both molecular and morphological purposes, but other fixation-preservation strategies were found to be preferable.

Specimens were deposited in the Scripps Institution of Oceanography Benthic Invertebrate Collection (**SIO-BIC**) and the Museo de Zoología, Universidad de Costa Rica (**MZUCR**, invertebrate collections; **UCR**, fish collection).

Taxonomic scope

We targeted benthic invertebrate macrofauna (retained on a 300 µm mesh and typically > 1 mm in size) and megafauna due to the nature of our sampling gear, although a few opportunistically collected exceptions such as large nematodes are reported. The meiofaunal (Neira et al. 2018; Gracia C et al. 2020), fungal (Rojas-Jimenez et al. 2020), and microbial communities of the Costa Rica seeps (Bailey et al. 2011; Niemann et al. 2013; Tavormina et al. 2013; Dekas et al. 2014; Case et al. 2015; Pasulka et al. 2017; Brzechffa and Goffredi 2021; Metcalfe et al. 2021) have been explored in other studies. Our sampling was not intended to be quantitative, as related investigations have explored macrofaunal species richness, biomass, density, and other ecological metrics at these seeps (Levin et al. 2012, 2015; Ashford et al. 2021b, 2021a; Pereira et al. 2021, 2022).

We reported only taxa that were linked to physical specimens, given the importance of museum vouchers for genetic characterization, species descriptions, scientific reproducibility, and the principles of Findable, Accessible, Interoperable, and Reusable (FAIR) research (Howell et al. 2020). Nonetheless, these expeditions also captured images and video of additional fauna (e.g., medusae, cephalopods, and fishes) that were not collected and that may form the basis of future studies.

Table 1. Submersible dives and collection events. AD = HOV Alvin, R/V Atlantis. S = ROV SuBastian, R/V Falkor. MC = multi-core. Multicores and plankton tows were deployed from R/V Atlantis. Coordinates reflect dive summaries as reported in the HOV Alvin dive logs (<https://ndsf.whoi.edu/data/>) and the R/V Falkor FK190106 cruise report. Dates reflect local time. Depth ranges reflect the minimum and maximum depth for sample collection events. In most cases, more precise coordinates and depths for specific animals are available on the SIO-BIC online database (<https://sioapps.ucsd.edu/collections/bi/>).

Cruise	Dive or deployment	Locality	Date	Latitude, Longitude	Depth, m
AT15-44	AD4501	Mound 12	2009-02-22	8.9300, -84.3135	984–997
	AD4502	Mound 12	2009-02-23	8.9285, -84.3132	987–997
	AD4503	Mound 12	2009-02-24	8.9308, -84.3072	967–995
	AD4504	Mound 11	2009-02-25	8.9208, -84.3054	1004–1011
	AD4505	Mound 11	2009-02-26	8.9198, -84.3055	1019–1025
	AD4506	Parrita Seep	2009-02-27	8.9718, -84.6235	1030–1179
	AD4507	Parrita Scar	2009-02-28	8.9353, -84.6465	1659–1667
	AD4508	Parrita Seep	2009-03-01	9.0303, -84.6230	1401–1419
	AD4509	Jacó Scar and Jacó Slope	2009-03-03	9.1172, -84.8425	974–1856
	AD4510	Jacó Summit	2009-03-04	9.1723, -84.7987	741–744
	AD4511	Mound 12	2009-03-05	8.9305, -84.3123	988–997
	AD4512	Quepos Slide	2009-03-06	8.8536, -84.2181	344–411
	AD4513	Jacó Scar	2009-03-07	9.1167, -84.8351	1744–1818
	MC1-2	Transition site near Mound 12 (~ 500 m from active seep)	2009-02-21	8.9316, -84.3168	1019
AT15-59	AD4586	Mound 12	2010-01-07	8.9308, -84.3130	982–998
	AD4587	Mound 12	2010-01-08	8.9306, -84.3123	990–996
	AD4588	Mound 12	2010-01-09	8.9308, -84.3125	995–997
	AD4589	Mound 12	2010-01-10	8.9298, -84.3121	997
	AD4590	Jacó Scar	2010-01-11	9.1176, -84.8395	1791–1800
	AD4591	Jacó Scar	2010-01-12	9.1182, -84.8391	1752–1795
	MC1	Transition site near Mound 12 (~ 400 m from active seep)	2010-01-06	8.9325, -84.3158	995
	MC4	Transition site near Mound 11 (~ 400 m from active seep)	2010-01-10	8.9208, -84.3016	1031
	Plankton Tow 6	Jacó Summit	2010-01-12	9.1713, -84.7987	0–350 (350 m wire out)
AT37-13	AD4906	Mound 12	2017-05-21	8.9308, -84.3128	995–1002
	AD4907	Mound 12	2017-05-22	8.9304, -84.3128	990–999
	AD4908	Mound 12	2017-05-23	8.9304, -84.3126	989–1001
	AD4909	Mound 12	2017-05-24	8.9305, -84.3125	967–1000
	AD4910	Mound 12	2017-05-25	8.9304, -84.3126	988–1004
	AD4911	Jacó Scar	2017-05-26	9.1151, -84.8468	1757–1892
	AD4912	Jacó Scar	2017-05-27	9.1154, -84.8362	1795–1859
	AD4913	Jacó Scar	2017-05-28	9.1156, -84.8401	1798–1908
	AD4914	Jacó Scar	2017-05-29	9.1175, -84.8395	1632–1886
	AD4915	Jacó Scar	2017-05-30	9.1180, -84.8404	1741–1885
	AD4916	Jacó Scar	2017-05-31	9.1193, -84.8428	1604–1854
	AD4917	Mound 12	2017-06-01	8.9293, -84.3150	965–1000
	AD4918	Quepos Slide	2017-06-02	8.8535, -84.2177	333–408
	AD4919	Quepos Slide	2017-06-03	8.8527, -84.2174	379–410
	AD4921	Quepos Slide	2017-06-04	8.8532, -84.2155	345–394
	AD4922	Mound 12	2017-06-05	8.9296, -84.3078	964–1009
	AD4923	Parrita Seep	2017-06-06	8.9759, -84.6238	1037–1097
	AD4924	Parrita Seep	2017-06-07	9.0305, -84.6202	1400–1410
AT42-03	AD4971	Jacó Scar	2018-10-17	9.1170, -84.8426	1746–1824
	AD4972	Jacó Scar	2018-10-18	9.1164, -84.8403	1746–1845
	AD4973	Jacó Scar	2018-10-19	9.1148, -84.8398	1784–1887
	AD4974	Mound 12	2018-10-20	8.9297, -84.3078	990–1010
	AD4975	Mound 12	2018-10-21	8.9310, -84.3075	988–1002
	AD4976	Jacó Scar	2018-10-22	9.1139, -84.8401	1836–1887
	AD4977	Jacó Scar	2018-10-23	9.1163, -84.8418	1783
	AD4978	Mound 12	2018-10-24	8.9294, -84.3143	996–999

Cruise	Dive or deployment	Locality	Date	Latitude, Longitude	Depth, m
AT42-03	AD4979	Quepos Slide	2018-10-25	8.8539, -84.2178	380–397
	AD4984	Mound 12	2018-10-30	8.9300, -84.3137	964–998
	AD4985	Mound 12	2018-10-31	8.9303, -84.3129	991–1002
	AD4986	Quepos Slide	2018-11-01	8.8540, -84.2195	308–379
	AD4987	Mound 12 West	2018-11-02	8.9292, -84.3167	995–1012
	AD4988	Mound 11	2018-11-03	8.9193, -84.3027	998–1025
	AD4989	Jacó Scar	2018-11-04	9.1174, -84.8417	1758–1792
	AD4990	Parrita Seep	2018-11-05	9.0321, -84.6197	1400–1435
FK190106	S0212	Jacó Scar	2019-01-06	9.1175, -84.8393	1780–1869
	S0213	Jacó Summit	2019-01-06	9.1734, -84.8038	730–820
	S0214	Jacó Scar	2019-01-07	9.1175, -84.8393	1780–1875
	S0215	Mound 12	2019-01-08	8.9307, -84.3126	982–1016
	S0216	Quepos Slide	2019-01-09	8.8539, -84.2193	275–404
	S0217	The Thumb	2019-01-10	9.0486, -84.3945	940–1074
	S0218	Parrita Scar	2019-01-11	8.9498, -84.6381	1110–1988
	S0219	Rio Bongo Scar	2019-01-13	9.2862, -85.2757	480–661
	S0220	Subduction Plume	2019-01-14	8.8785, -84.8695	3399–3601
	S0230	Mound Jaguar	2019-01-25	9.6558, -85.8813	1895–2000

Specimen identification

Following preliminary morphological identification during shipboard processing, specimens were identified to the lowest possible taxonomic level based on genetics and/or morphology. Genetic identification was facilitated by querying sequences against the NCBI GenBank database (Clark et al. 2016) using the nucleotide BLAST (blastn) suite (<https://blast.ncbi.nlm.nih.gov/>) (Zhang et al. 2000; Boratyn et al. 2013). Biogeography assessments and access to taxonomic literature were facilitated by the Global Biodiversity Information Facility (GBIF) (GBIF: The Global Biodiversity Information Facility 2024) and the Biodiversity Heritage Library (Gwinn and Rinaldo 2009).

Taxonomic uncertainty was expressed using recommended terminology and practices for open nomenclature (Sigovini et al. 2016; Horton et al. 2021). For example, the abbreviation “stet.” (*stet.*) is used to indicate that further identification to a lower taxonomic level was not attempted given the limitations of the specimen or of available taxonomic resources. Uncertainty at a given taxonomic level is indicated by the abbreviation “inc.” (*incerta, incertus, incertum*). Undescribed species that have been identified with certainty as new to science but have not yet been formally described or referenced in the literature, are notated with a unique alphanumeric code linked to a museum voucher, e.g., “sp. SIO_BIC_A00001” as opposed to a potentially non-unique designation such as “sp. 1”. Some morphospecies are also notated with unique voucher-linked alphanumeric codes, when multiple morphospecies must be distinguished from one another but cannot be matched with certainty to known species.

DNA extraction and sequencing

Genomic DNA was extracted following the manufacturer’s protocol for commercial kits such as the DNeasy Tissue Kit (Qiagen); the EZNA Micro-Elute Genomic DNA Kit (Omega Bio-Tek); or the *Quick*-DNA Miniprep, Microprep Plus, or 96 Plus Kit (Zymo Research, Irvine, CA and Tustin, CA). Polymerase chain reaction (PCR) amplification of phylogenetically informative gene fragments was performed using the primer pairs summarized in Table 2. A typical PCR included 1 µl of each

Table 2. PCR primers and temperature profiles. Mitochondrial genes: COI = cytochrome c oxidase subunit I; COIII = cytochrome c oxidase subunit III; 16S = ribosomal RNA 16S subunit. Nuclear genes: 18S = ribosomal RNA 18S subunit.

Amplified gene fragment	Primer pair	References for primer sequences	Temperature profile	Taxa
COI	LCO1490/ HCO2198	Folmer et al. 1994	94 °C/180s – (94 °C/30s – 47 °C/45s – 72 °C/60s) * 5 cycles – (94 °C/30s – 52 °C/45s – 72 °C/60s) * 30 cycles – 72 °C/300s	Annelida, Arthropoda, Mollusca, Nemertea
COI	dgLCO/ dgHCO	Meyer 2003	95 °C/120s – (95 °C/40s – 45 °C/40s – 72 °C/60s) * 35 cycles – 72 °C/420s or 95 °C/300s – (95 °C/30s – 48 °C/30s – 72 °C/45s) * 35 cycles – 72 °C/300s	Annelida, Mollusca, Holothuroidea, Caridea
COI	PolyLCO/ PolyHCO	Carr et al. 2011	95 °C/180 s – (95 °C/40 s – 42 °C/40 s – 72 °C/50 s) * 40 cycles – 72 °C/300 s	Annelida
COI	HCO2198/LCO_Apl	Folmer et al. 1994; Bergmeier et al. 2019	95 °C/60s – (95 °C/20s – 52 °C/15s – 72 °C/30s) * 40 cycles – 72 °C/420s	Aplacophora
COI	HCO2198/ CrustF2	Costa et al. 2007	95 °C/60s – (95 °C/30s – 42 °C/90s – 72 °C/60s) * 35 cycles – 72 °C/300s	Arthropoda
COI	COIceF/ COIceR	Hoareau and Boissin 2010	95 °C/180s – (94 °C/45s – 48 °C/70s – 72 °C/80s) * 40 cycles – 72 °C/600s	Echinodermata
COI	ECOLa/ HCO2198	Folmer et al. 1994; Knott and Wray 2000 combined as in Deagle et al. 2003	94 °C/240s – (94 °C/30s – 50 °C/30s – 72 °C/45s) * 35 cycles – 72 °C/300s	Astroidea
COI	Fsco1/ Co13r	Helgen and Rouse 2006	94 °C/180s – (94 °C/45s – 48 °C/45s – 72 °C/60s) * 35 cycles – 72 °C/480s	Crinoidea
COI	COlef/ COler	Arndt et al. 1996	95 °C/120s – (95 °C/30s – 48 °C/30s – 72 °C/45s) * 35 cycles – 72 °C/600s	Holothuroidea, Echinoidea
COI	VesLCO/ VesHCO	Peek et al. 1997	94 °C/240s – (94 °C/40s – 40 °C/40s – 72 °C/60 s) * 40 cycles – 72 °C/600 s	Vesicomysidae
COI	jgLCO1490/ jgHCO2198	Geller et al. 2013	95 °C/300s – (95 °C/30s – 48 °C/30s – 72 °C/45s) * 35 cycles – 72 °C/300s	<i>Hyalogyrina</i>
COIII	COIIIF/ COIIIR	Geller and Walton 2001	95 °C/120s – (95 °C/30s – 45 °C/30s – 72 °C/60s) * 30 cycles – 72 °C/300s	Actiniaria
16S	16SarL/ 16SbrH	Palumbi 1996; Palumbi et al. 2002	95 °C/180s – (95 °C/40s – 50 °C/40s – 68 °C/50 s) * 35 cycles – 68 °C/300 s or 95 °C/180s – (95 °C/40s – 50 °C/40s – 72 °C/50 s) * 40 cycles – 72 °C/300 s	Annelida, Nemertea, Holothuroidea, Polyplacophora, <i>Paracrangon areolata</i> , <i>Grimothea monodon</i>
16S	ANEM16SA/ ANEM16SB	Geller and Walton 2001	95 °C/120s – (95 °C/30s – 60 °C/30s – 72 °C/60s) * 30 cycles – 72 °C/300s	Anthozoa
16S	16S_arL_solenos (CGACTGTTTAACAAAAACATTGCTC)/ 16S_brH_solenos (CCGATTGAAGTACATCATGTAG)	this work (K. Kocot)	95 °C/60s – (95 °C/20s – 52 °C/15s – 72 °C/30s) * 40 cycles – 72 °C/420s	Aplacophora
16S	AnnF/ 16Sb	Edgecombe et al. 2002; Sjölin et al. 2005 combined as in Stiller et al. 2013	94 °C/120s – (94 °C/40s – 60 °C/40s – 70 °C/45s) * 35 cycles – 72 °C/420s	Sabellidae, Macellicephalinae
18S	Three overlapping fragments: 1F/5R, 3F/bi, a2.0/9R	Giribet et al. 1996, 1999 combined as in Stiller et al. 2013	1F/5R: 95 °C/180s – (95 °C/60s – 49 °C/30s – 72 °C/90s) * 40 cycles – 72 °C/480s; 3F/bi: 95 °C/180s – (95 °C/30s – 52 °C/30s – 72 °C/90s) * 40 cycles – 72 °C/480s; a2.0/9R: 95 °C/180s – (95 °C/30s – 49 °C/30s – 72 °C/90s) * 40 cycles – 72 °C/480s	Sabellidae
18S	Three overlapping fragments: TimA/1100R2, 3F/bi, a2.0/9R	Giribet et al. 1996, 1999; Norén and Jondelius 1999 combined as in Stiller et al. 2013	TimA/1100R2: 94 °C/180s – (94 °C/30s – 53 °C/45s – 72 °C/120s) * 40 cycles – 72 °C/300s; 3F/bi, a2.0/9R: see previous	Sabellidae
18S	Sol18F/ Sol18R	Neulinger et al. 2006	94 °C/300s – (91 °C/40s – 50 °C/40s – 72 °C/90s) * 40 cycles – 72 °C/300s	Solemyidae

primer (10 μ M), 2 μ l of genomic DNA, and the appropriate concentration of a commercially available reagents such as Apex 2.0x Taq Red DNA Polymerase Master Mix (Genesee Scientific), Hot Start Taq PCR Master Mix 2X (VWR), or Conquest PCR Master Mix (Lamda Biotech, St. Louis, MO). PCR products were purified with ExoSAP-IT (USB Corporation, Cleveland, OH) or the EZNA Cycle Pure Kit (Omega Bio-Tek). Sanger sequencing was performed by Eurofins Genomics (Louisville, KY), GeneWiz (South Plainfield, NJ), or Retrogen, Inc. (San Diego, CA). Consensus sequences were assembled using Geneious (<https://www.geneious.com>).

Haplotype networks

Sequences were aligned using the MAFFT online service v. 7.471, option L-INS-I (Katoh et al. 2018). Haplotype networks were created with PopART v. 1.7 (Leigh and Bryant 2015) using the TCS algorithm (Clement et al. 2002).

Scanning electron microscopy (SEM)

Selected aplacophoran specimens were dried and mounted on stubs without critical point drying or sputter coating. Specimens were imaged on a Phenom Pro SEM.

Permits

Specimen collection and field operations were performed under the following permits issued by CONAGEBIO (Comisión Nacional para la Gestión de la Biodiversidad), INCOPECA (Instituto Costarricense de Pesca y Acuicultura), and SINAC (Sistema Nacional de Áreas de Conservación) under MINAE (Ministerio de Ambiente y Energía), Government of Costa Rica: INCOPECA-CPI-003-12-2018, R-070-2018-OT-CONAGEBIO, SINAC-CUSBSE-PI-R-032-2018, SINAC-SE-CUS-PI-R-035-2017. In accordance with the Nagoya Protocol on Access and Benefit Sharing, DNA sequencing for this project was authorized by the Contract for the Grant of Prior Informed Consent between MINAE-SINAC-ACMC and Jorge Cortés-Núñez for the Basic Research Project: “FK190106-Cuantificación de los vínculos biológicos, químicos y físicos entre las comunidades quimiosintéticas con el mar profundo circundante.”

Taxonomic listing

For each taxonomic entry, we summarize the known localities and depths of occurrences at the CRM seeps, incorporating both published references and additional material examined in this work. Representative voucher specimens and their associated DNA sequences are listed by dive/deployment number (details in Table 1). Catalog numbers indicate a morphological voucher linked to a tissue sample suitable for genetic analysis, unless otherwise noted. For some groups such as octocorals, morphological vouchers and genetic tissue samples were deposited with separate institutions according to local expertise.

We also summarize the known localities and depths of occurrences beyond the CRM seeps. We indicate new biogeographic records, new depth records (defined here as at least 100 m from a previously reported minimum or maximum depth), and new seep records of species previously associated with vents

or organic falls. Exact collection depths are provided where possible; approximations are indicated by ~ .

References indicate previously published records from the CRM seeps. Original species descriptions are indicated by **. References generally include voucher specimen listings with representative images and DNA sequences; where possible, we provide any missing components. Specimen catalog numbers pertain to SIO-BIC unless otherwise indicated. GenBank numbers refer to COI sequences unless otherwise indicated. New sequences are shown in bold.

For higher-level taxonomy, we follow the World Register of Marine Species (WoRMS Editorial Board 2024) or taxon-specific references. We notate higher classification according to the recommended best practice for the DarwinCore term higherClassification (Darwin Core Maintenance Group 2021), with full taxonomic authorities and Linnaean ranks available in the applicable references. Our ordering of major animal groups reflects the phylogeny in Dunn et al. (2014). Entries are listed alphabetically within each category.

Annelida

We list entries following the taxonomic arrangement in Rouse et al. (2022).

Annelida | Polychaeta | Errantia | Protodriliformia

Protodrilidae *stet.*

Fig. 3A

Material examined. AD4906: A8261; AD4923: A8456 ([PQ449314](#)).

Localities. Mound 12 (1002 m), Parrita Seep (~ 1040–1101 m).

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida |
Aphroditiformia | Polynoidae

***Bathykurila* sp. A** *sec. Glover et al. 2005*

Fig. 3B, C

Material examined. AD4906: A8264; S0213: A10050 ([PQ449253](#)); S0217: A10086.

Localities. Jacó Summit (~ 730–820 m), Mound 12 (~ 997–1002 m), Mound 11 (1004–1040 m), The Thumb (1072 m).

Remarks. COI sequences of this morphospecies were > 99.42% identical to those of *Bathykurila* haplotype group A (GenBank [DQ074778.1](#), [DQ074779.1](#), [DQ074780.1](#)), which morphologically resembles *B. guaymasensis* Pettibone, 1989 as discussed in Glover et al. (2005).

***Branchinotogluma* sp. SIO_BIC_A8265**

Fig. 3D, E

Material examined. AD4907: A8265 ([OR682087](#)).

Localities. Mound 12 (~ 990–999 m).

Remarks. An undescribed species. Specimen A8265 was associated with experimentally deployed wood.

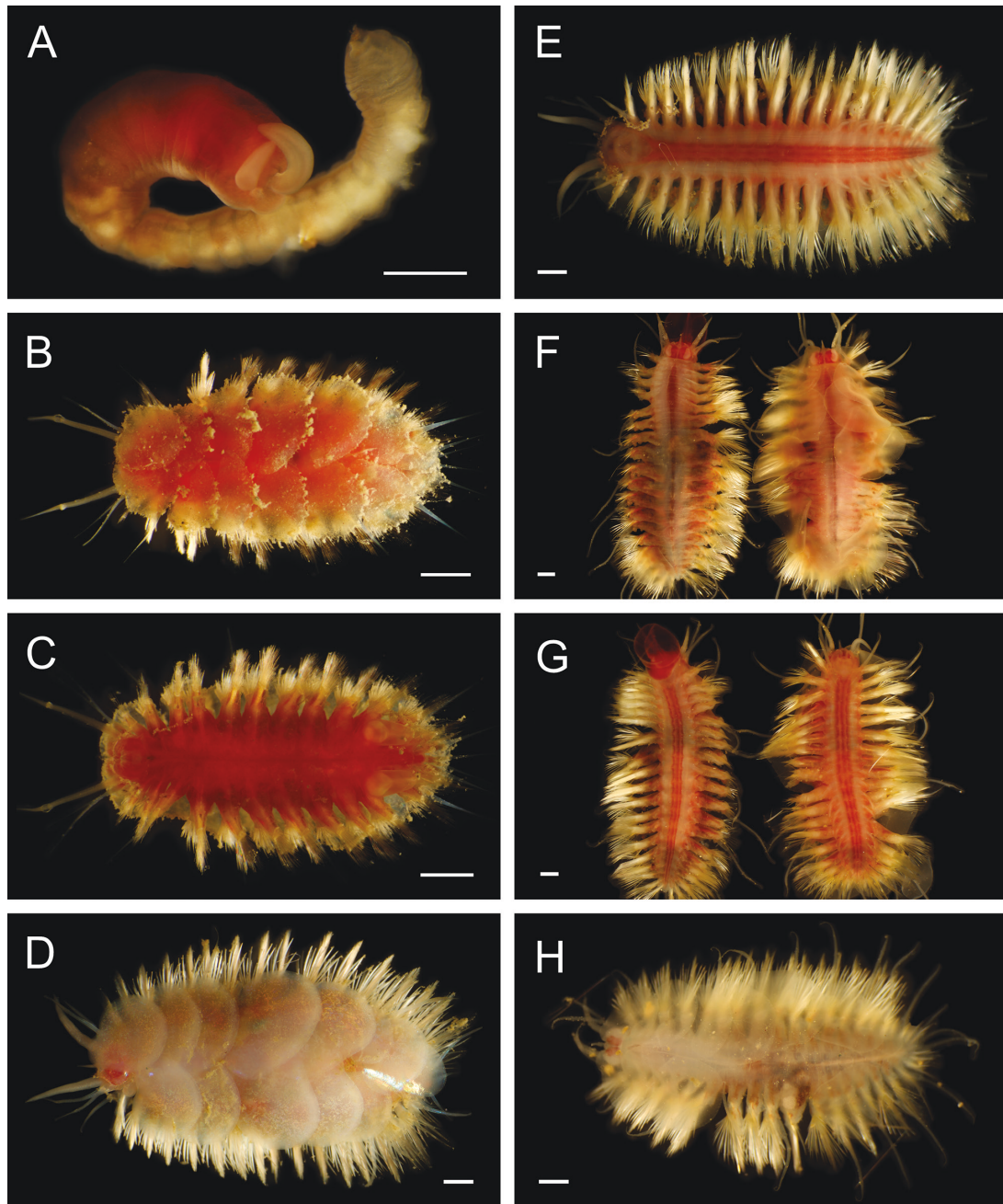


Figure 3. Annelida: Protodrilidae and Polynoidae, representative live images **A** Protodrilidae stet. (A8456) **B** *Bathykurila* sp. A sec. Glover et al. 2005 (A10050, dorsal view) **C** *Bathykurila* sp. A sec. Glover et al. 2005 (A10050, ventral view) **D** *Branchinotogluma* sp. SIO_BIC_A8265 (A8265, dorsal view) **E** *Branchinotogluma* sp. SIO_BIC_A8265 (A8265, ventral view) **F** *Branchinotogluma* sp. SIO_BIC_A8460 (A8460 and A16362, dorsal view) **G** *Branchinotogluma* sp. SIO_BIC_A8460 (A8460 and A16362, ventral view) **H** *Branchinotogluma* sp. SIO_BIC_A8460 (A8461, dorsal view). Scale bars: 1 mm.

***Branchinotogluma* sp. SIO_BIC_A8460**

Fig. 3F–H

Material examined. AD4913: A13252 ([OR682108](#)); AD4924: A8460 ([OR682111](#)), A8461, A16362; S0218: A10190 ([OR682109](#)).

Localities. Parrita Scar (1364 m), Parrita Seep (~ 1400–1410 m), Jacó Scar (1847 m).

Remarks. An undescribed species.

***Branchinotogluma* sp. SIO_BIC_A9682**

Fig. 4A–C

Material examined. AD4505: A1363 ([OR682006](#)); AD4924: A8459 ([OR682020](#)); AD4972: A9682 ([OR682056](#)); AD4978: A9763 ([OR682070](#)); S0230: A10185 ([OR682045](#)), A10187 ([OR682037](#)).

Localities. Mound 12 (~ 1000 m), Mound 11 (1025 m), Parrita Seep (~ 1400 m), Jacó Scar (~ 1800 m), Mound Jaguar (1908–1909 m).

Remarks. An undescribed species.

***Branchipolynoe eliseae* Lindgren, Hatch, Hourdez, Seid & Rouse, 2019**

Fig. 4D

Reference. Lindgren et al. 2019**.

Localities. Mound 12 (997 m; type locality), Jacó Scar (~ 1752–1800 m).

Distribution. Known only from the CRM seeps.

Remarks. Symbiont of the mussels *Bathymodiolus billschneideri* and *Ba. nancyschneiderae* (Lindgren et al. 2019), typically with one adult worm and sometimes several very small juvenile worms per mussel. Possibly also found in *Ba. earlougheri*, whose locality and depth ranges overlap with the two known host mussel species (McCowin et al. 2020). The absence of *B. eliseae* occurrences in *Ba. earlougheri* is likely an artefact of limited sample size (Lindgren et al. 2019).

***Branchipolynoe halliseyae* Lindgren, Hatch, Hourdez, Seid & Rouse, 2019**

Fig. 4E

Reference. Lindgren et al. 2019**.

Localities. Mound 12 (~ 1000 m; type locality), Parrita Seep (~ 1400 m), Jacó Scar (1758–1811 m).

Distribution. Known only from the CRM seeps.

Remarks. Symbiont of the mussels *Bathymodiolus billschneideri*, *Ba. nancyschneiderae*, and *Ba. earlougheri* (Lindgren et al. 2019), typically with one adult worm and sometimes several very small juvenile worms per mussel.

***Branchipolynoe kajsae* Lindgren, Hatch, Hourdez, Seid & Rouse, 2019**

Fig. 4F

Reference. Lindgren et al. 2019**.

Localities. Mound 12 (~ 1000 m; type locality), Parrita Seep (~ 1400 m), Jacó Scar (~ 1800 m).

Distribution. Known only from the CRM seeps.

Remarks. Symbiont of the mussels *Bathymodiolus billschneideri*, *Ba. nancyschneiderae*, and *Ba. earlougheri* (Lindgren et al. 2019), typically with one adult worm and sometimes several very small juvenile worms per mussel.

***Branchipolynoe meridae* Lindgren, Hatch, Hourdez, Seid & Rouse, 2019**

Fig. 4G

Reference. Lindgren et al. 2019**.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m; type locality).

Distribution. Known only from the CRM seeps.

Remarks. Symbiont of the mussels *Bathymodiolus billschneideri* and *Ba. earlougheri* (Lindgren et al. 2019), typically with one adult worm and sometimes several very small juvenile worms per mussel. Possibly also found in *Ba. nancyschneiderae*, which co-occurs with *Ba. earlougheri* at Mound 12 (McCowin et al. 2020). The absence of *B. eliseae* occurrences in *Ba. nancyschneiderae* is likely an artefact of limited sample size (Lindgren et al. 2019).

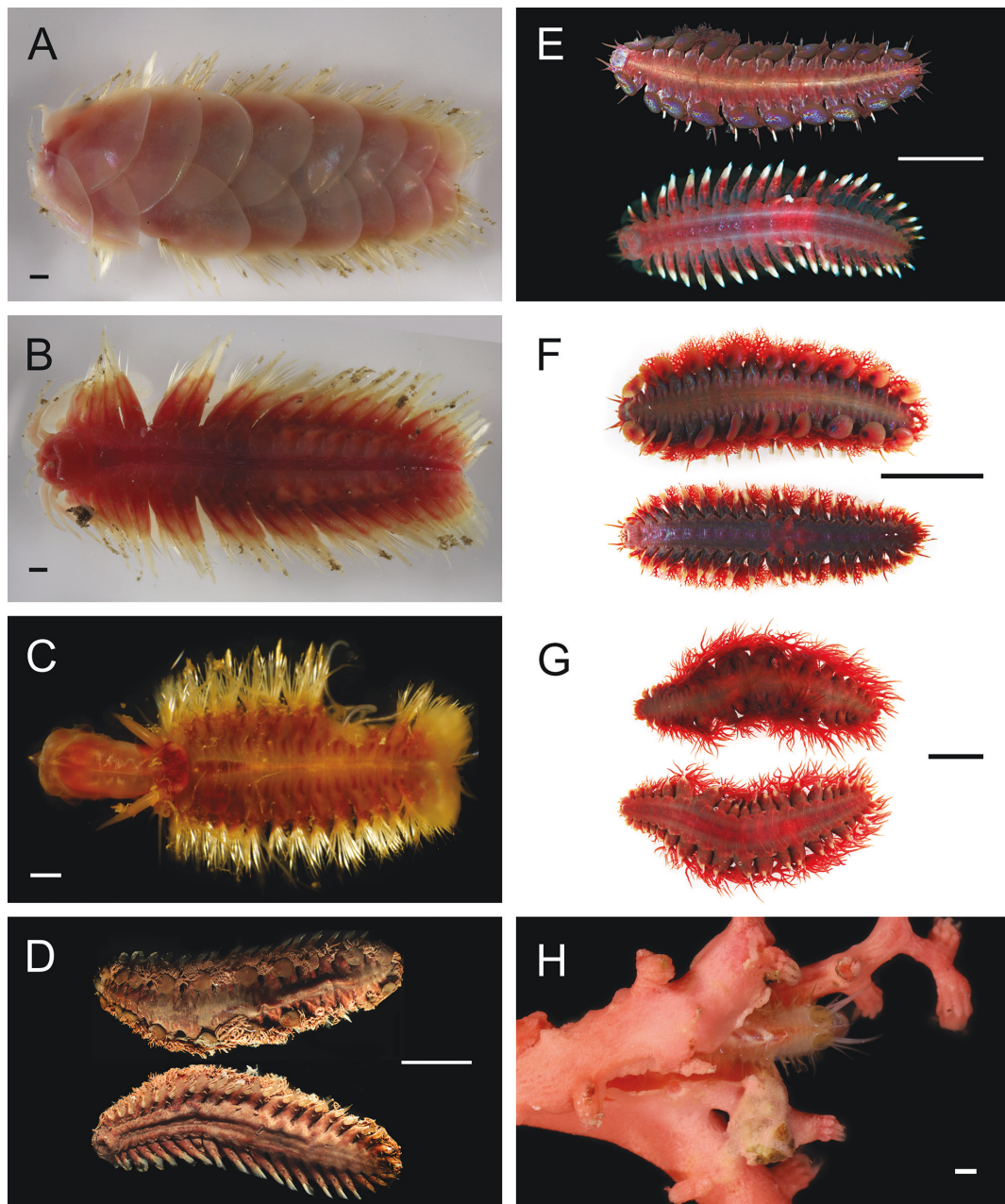


Figure 4. Annelida: Polynoidae, representative images. Live specimens are depicted unless otherwise specified **A** *Branchinotogluma* sp. SIO_BIC_A9682 (A9682, dorsal view) **B** *Branchinotogluma* sp. SIO_BIC_A9682 (A9682, ventral view) **C** *Branchinotogluma* sp. SIO_BIC_A9682 (A9763, dorsal view without scales) **D** *Branchipolynoe eliseae* (A6660, preserved specimen, dorsal and ventral views) **E** *Branchipolynoe halliseyae* (A1322, dorsal and ventral views) **F** *Branchipolynoe kajsae* (A6611, dorsal and ventral views) **G** *Branchipolynoe meridae* (A6616, dorsal and ventral views) **H** *Gorgoniapolynoe* cf. *caeciliae* (A8485, with host coralliid Co2947). Scale bars: 1 mm (**A–C, H**); 1 cm (**D–G**).

***Gorgoniapolynoe cf. caeciliae* (Fauvel, 1913)**

Figs 4H, 5A

Material examined. AD4506: A1549; AD4923: A8455 ([PQ449313](#)), A8485.

Localities. Parrita Seep (~ 1030–1094 m).

Remarks. Associated with coralliid octocorals: A1549 with coral Co2271, A8455 and A8485 with coral Co2947. The COI sequence of A8455 was 98.20% identical to a reference sequence of *Gorgoniapolynoe cf. caeciliae* molecular operational taxonomic unit (MOTU) 1 from the central Atlantic ([ON479554.1](#)), representing one of two lineages in a potential cryptic species complex with inter-lineage COI distances of 2–7% (Maxwell et al. 2022). The CRM specimens warrant further detailed comparison to these Atlantic *G. cf. caeciliae* lineages as well as to the eastern Pacific species *G. guadalupensis* Pettibone, 1991, which currently has no available reference sequences. *Gorgoniapolynoe guadalupensis* was originally described in association with *Hemicorallium imperiale* (Bayer, 1955) off Guadelupe Island, western Mexico, 1000–2000 m, and has been recorded from seamounts in the eastern Pacific (Fieberling Guyot, off southern California) and central Pacific (Markus Nekar Chain, west of the Hawaiian Islands) to a minimum known depth of 600 m (Pettibone 1991).

***Macellicephala* sp. SIO_BIC_A8368**

Fig. 5B

Material examined. AD4913: A8368 ([PQ449306](#)).

Localities. Jacó Scar (~ 1817–1896 m).

***Macellicephala* sp. SIO_BIC_A9775**

Fig. 5C

Material examined. AD4975: A9775 ([PQ449325](#)).

Localities. Mound 12 (1000 m).

Remarks. An undescribed species. At least six individuals were associated with the ambulacral groove of an asteroid, *Thrissacanthias penicillatus* (E7246).

***Macellicephala* sp. SIO_BIC_A10055**

Fig. 5D, E

Material examined. S0213: A10055 ([OP648305](#)).

Localities. Jacó Summit (~ 730–820 m).

***Macellicephala* sp. SIO_BIC_A10094**

Fig. 5F, G

Material examined. S0219: A10094 ([PQ449266](#)).

Localities. Rio Bongo Scar (659 m).

Remarks. Afflicted with copepod parasites.

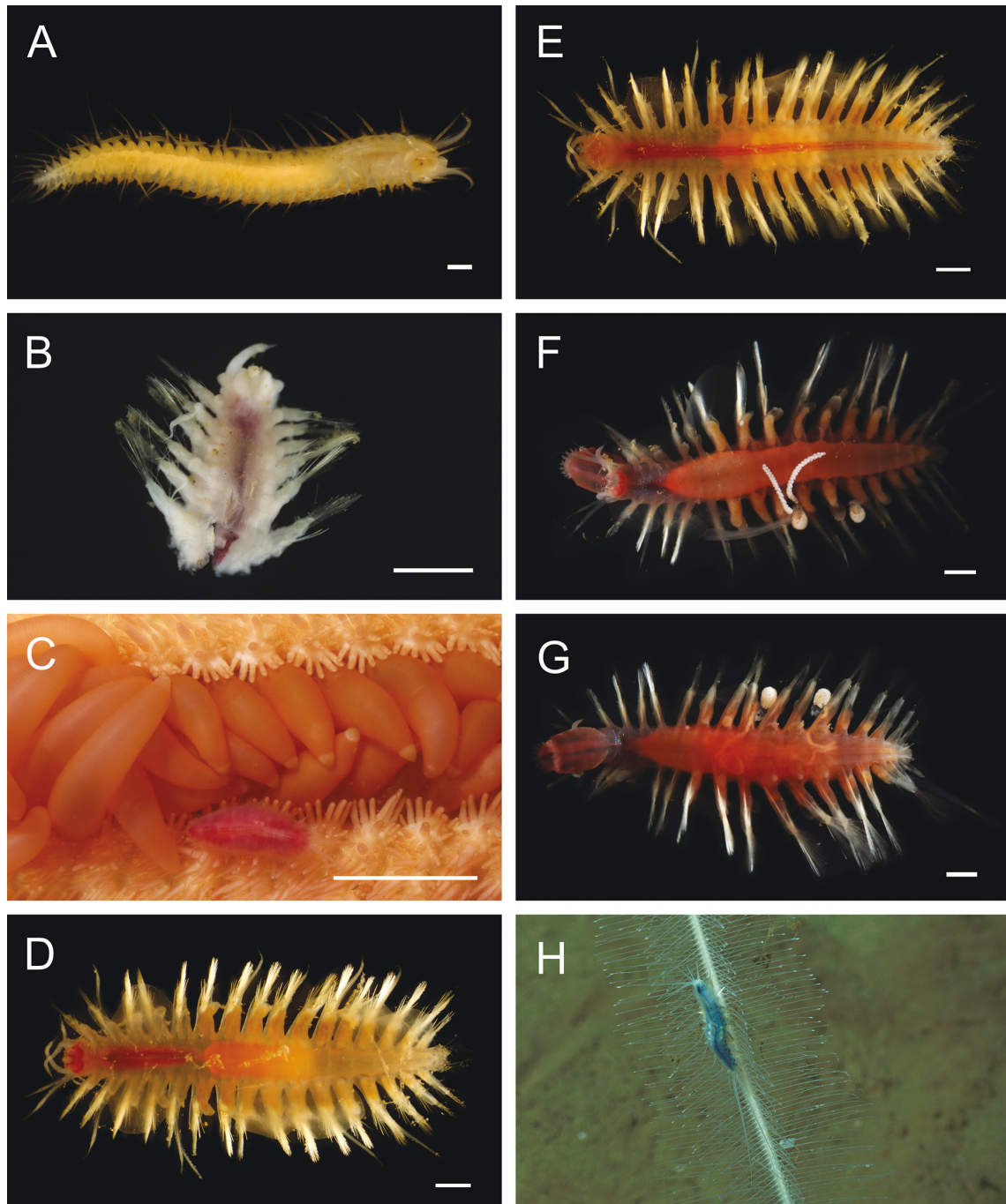


Figure 5. Annelida: Polynoidae, representative images. Live specimens are depicted unless otherwise specified **A** *Gorgoniapolynoe* cf. *caeciliae* (A8485, removed from host Co2947) **B** *Macellicephala* sp. SIO_BIC_A8368 (A8368, preserved specimen) **C** *Macellicephala* sp. SIO_BIC_A9775 (A9775) **D** *Macellicephala* sp. SIO_BIC_A10055 (A10055, dorsal view) **E** *Macellicephala* sp. SIO_BIC_A10055 (A10055, ventral view) **F** *Macellicephala* sp. SIO_BIC_A10094 (A10094, dorsal view) **G** *Macellicephala* sp. SIO_BIC_A10094 (A10094, ventral view) **H** *Macellicephala* sp. SIO_BIC_A10099 (A10099, *in situ* on host cladorhizid sponge P1754). Credit: ROV SuBastian/Schmidt Ocean Institute. Scale bars: 1 mm (**A**, **B**, **D–G**); 1 cm (**C**).

***Macellicephala* sp. SIO_BIC_A10099**

Figs 5H, 6A

Material examined. S0220: A10099 (OP648306; 16S: PQ304650).

Localities. Subduction Plume (3601 m).

Remarks. An undescribed species associated with a cladorhizid sponge, P1754.

Macellicephalinae sp. SIO_BIC_A8365

Fig. 6B, C

Material examined. AD4914: A8365 ([PQ449304](#)).

Localities. Jacó Scar (1839 m).

Remarks. Associated with a holothuroid, *Achlyonice* stet. (E7042 or E7043).

Macellicephalinae sp. SIO_BIC_A8458

Fig. 6D

Material examined. AD4923: A8458 ([PQ449315](#)).

Localities. Parrita Seep (~ 1037–1108 m).

Macellicephalinae sp. SIO_BIC_A10186

Fig. 6E, F

Material examined. S0230: A10186.

Localities. Mound Jaguar (1909 m).

***Peinaleopolynoe elvisi* Hatch, Liew, Hourdez & Rouse, 2020**

Fig. 6G

Reference. Hatch et al. 2020**.

Additional material examined. S0214: A10059 ([PQ449258](#)).

Localities. Jacó Scar (1845–1887 m).

Distribution. Also known from a whale fall in Monterey Submarine Canyon, off California, 1820 m (type locality) and from cow bones experimentally deployed at 2091 m at Seamount 1, which lies on the CRM ca 41 km southwest of Jacó Scar (Hatch et al. 2020).

Remarks. A10059 was associated with a naturally occurring wood fall. All known occurrences of *P. elvisi* have been associated with vertebrate bones or wood (naturally occurring and experimentally deployed, for both substrates) (Hatch et al. 2020).

***Peinaleopolynoe mineoi* Hatch, Liew, Hourdez & Rouse, 2020**

Fig. 6H

Reference. Hatch et al. 2020**.

Localities. Mound 12 (992–1011 m; type locality), Mound 11 (1010 m).

Distribution. Known only from the CRM seeps.

Remarks. All known occurrences of *P. mineoi* have been associated with wood (naturally occurring and experimentally deployed) or experimentally deployed vertebrate bones (Hatch et al. 2020).

Polynoidae sp. SIO_BIC_A8426

Fig. 7A–C

Material examined. AD4922: A8426 ([PQ449308](#)).

Localities. Mound 12 (1006 m).

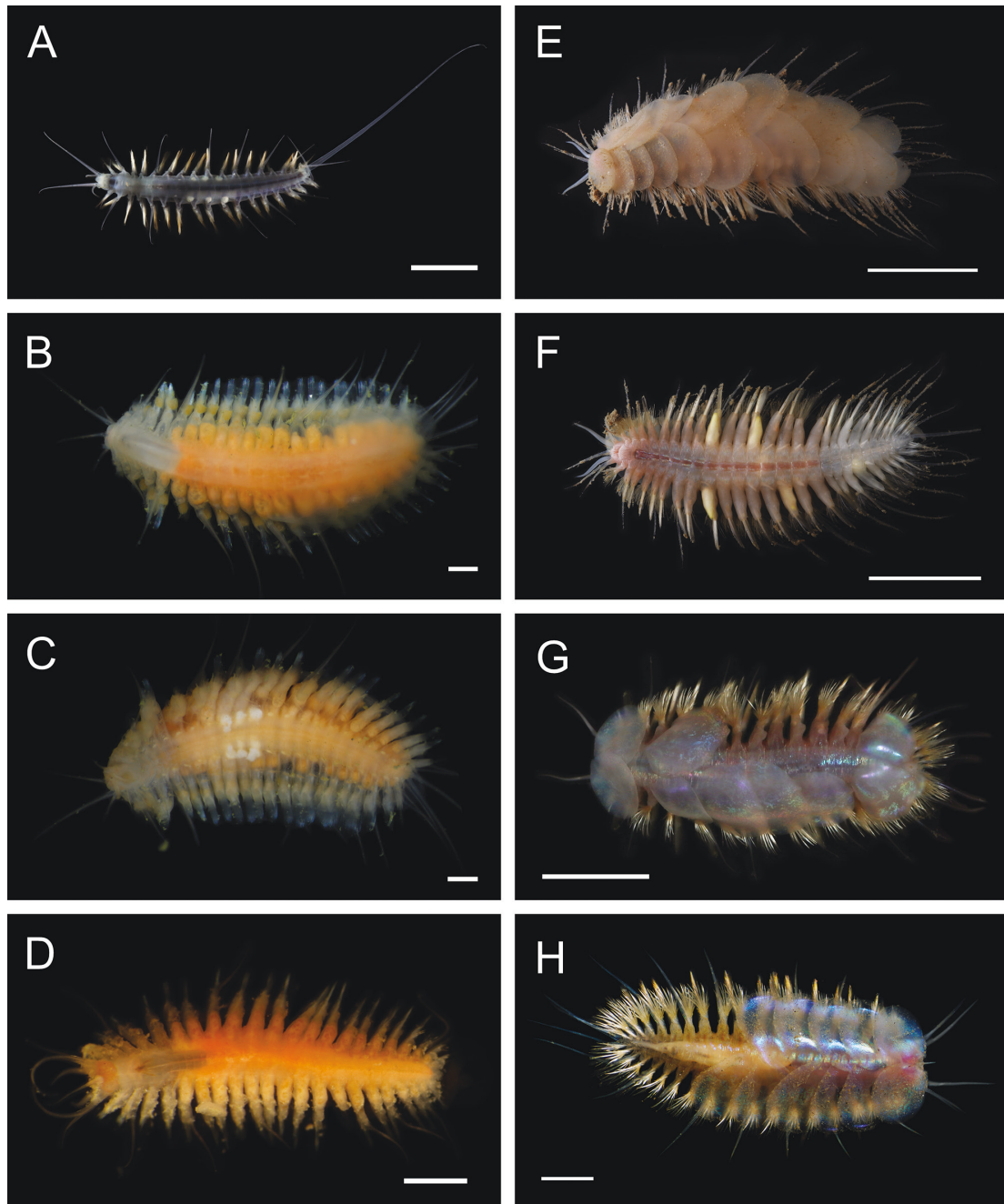


Figure 6. Annelida: Polynoidae, representative live images **A** *Macellicephala* sp. SIO_BIC_A10099 (A10099) **B** *Macellicephalinae* sp. SIO_BIC_A8365 (A8365, dorsal view) **C** *Macellicephalinae* sp. SIO_BIC_A8365 (A8365, ventral view) **D** *Macellicephalinae* sp. SIO_BIC_A8458 (A8458) **E** *Macellicephalinae* sp. SIO_BIC_A10186 (A10186, dorsal view) **F** *Macellicephalinae* sp. SIO_BIC_A10186 (A10186, ventral view) **G** *Peinaleopolynoe elvisi* (A10059) **H** *Peinaleopolynoe mineoi* (MZUCR 1001-01). Scale bars: 1 cm (**A**, **G**); 1 mm (**B–F**, **H**).

Remarks. Associated with a holothuroid, *Bathyplores* sp. SIO_BIC_E7063.

Polynoidae sp. SIO_BIC_A9652

Fig. 7D, E

Material examined. AD4972: A9720; AD4973: A9652 ([PQ449319](#)), A9653.

Localities. Jacó Scar (1784 m).

Polynoidae sp. SIO_BIC_A9714

Fig. 7F, G

Material examined. AD4974: A9714 ([PQ449321](#)).

Localities. Mound 12 (~ 1001–1003 m).

Polynoidae sp. SIO_BIC_A10082

Figs 7H, 8A

Material examined. AD4919: A8421; AD4986: A9899; S0216: A10082 ([PQ449262](#)).

Localities. Quepos Slide (~ 308–410 m)

Remarks. A10082 was associated with bones from a naturally occurring sailfish carcass.

Polynoidae sp. SIO_BIC_A10096

Fig. 8B, C

Material examined. S0219: A10095, A10096 ([PQ449267](#)), A10098, A10109, A10110.

Localities. Rio Bongo Scar (~ 480–650 m).

Remarks. A10096 and A10098 were commensals in a sponge, *Farrea occa* (P1753).

Polynoidae sp. SIO_BIC_A10189

Fig. 8D, E

Material examined. AD4913: A8369; S0230: A10189 ([PQ449271](#)).

Localities. Jacó Scar (~ 1817–1896 m), Mound Jaguar (2000 m).

Remarks. A10189 was associated with a naturally occurring wood fall.

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida |
Aphroditiformia | Syllidae

Anguillosyllis sp. SIO_BIC_A9613

Material examined. AD4975: A9613; AD4978: A9771.

Localities. Mound 12 (~ 992–999 m).

Remarks. An undescribed species. A9613 was associated with experimentally deployed wood and bone at 992 m.

Anguillosyllis sp. SIO_BIC_A12403

Fig. 8F

Reference. (Aguado et al. 2012), in which the published DNA sequences ([JF903571](#), [JF903680](#), [JF903756](#)) correspond to specimen A12403 from AT15-59, MC1.

Localities. Near Mound 12 (995 m).

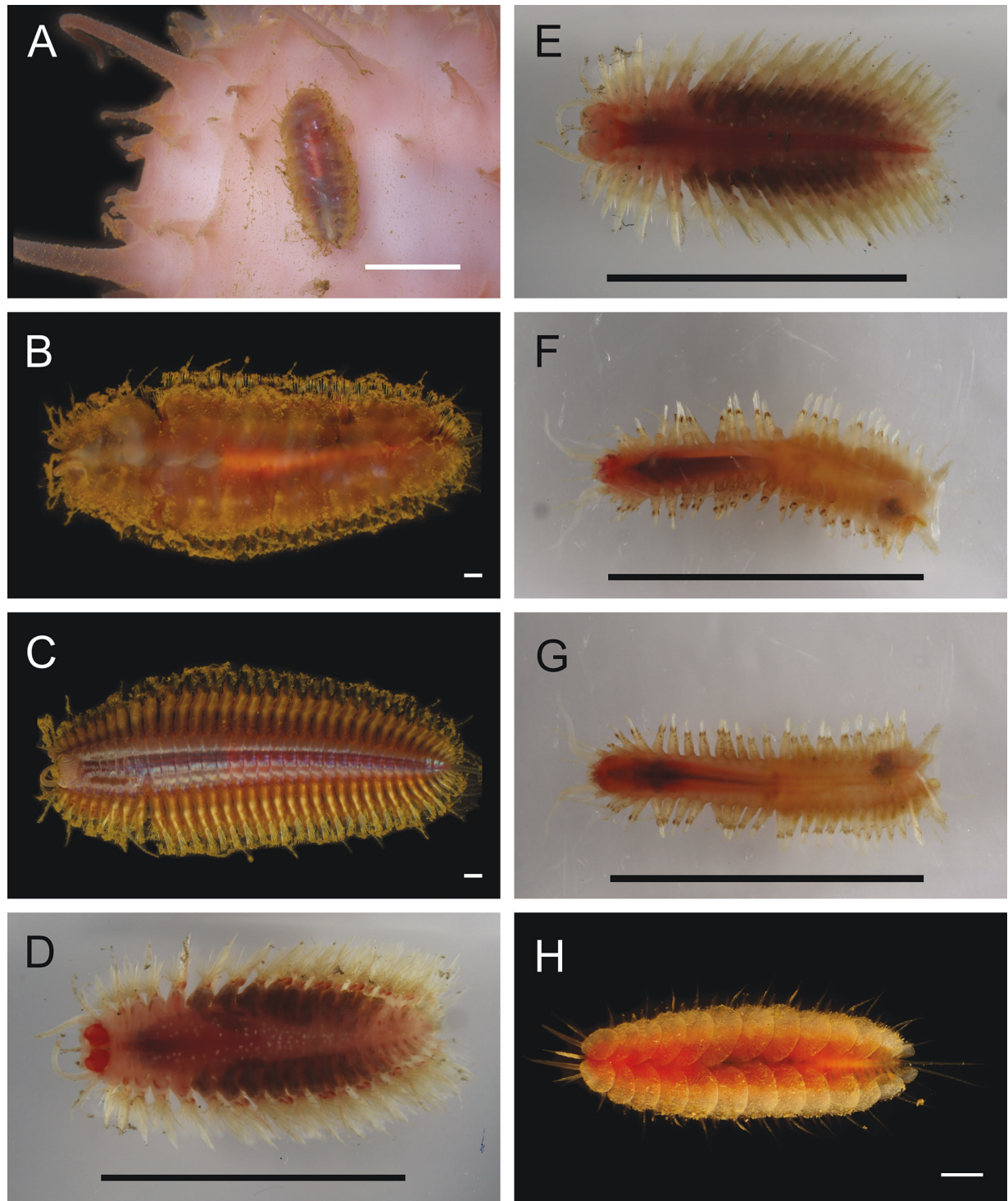


Figure 7. Annelida: Polynoidae, representative live images **A** Polynoidae sp. SIO_BIC_A8426 (A8426, on host holothuroid E7063) **B** Polynoidae sp. SIO_BIC_A8426 (A8426, dorsal view) **C** Polynoidae sp. SIO_BIC_A8426 (A8426, ventral view) **D** Polynoidae sp. SIO_BIC_A9652 (A9652, dorsal view) **E** Polynoidae sp. SIO_BIC_A9652 (A9652, ventral view) **F** Polynoidae sp. SIO_BIC_A9714 (A9714, dorsal view) **G** Polynoidae sp. SIO_BIC_A9714 (A9714, ventral view) **H** Polynoidae sp. SIO_BIC_A10082 (A10082, dorsal view). Scale bars: 1 cm (**A**, **D–G**); 1 mm (**B**, **C**, **H**).

Remarks. An undescribed species, morphologically similar to *Anguillosyllis* sp. SIO_BIC_A9613. This specimen was collected in a sediment core adjacent to Mound 12, ca 400 m from known sites of active seepage and likely representing the far-transition zone to the surrounding environment.

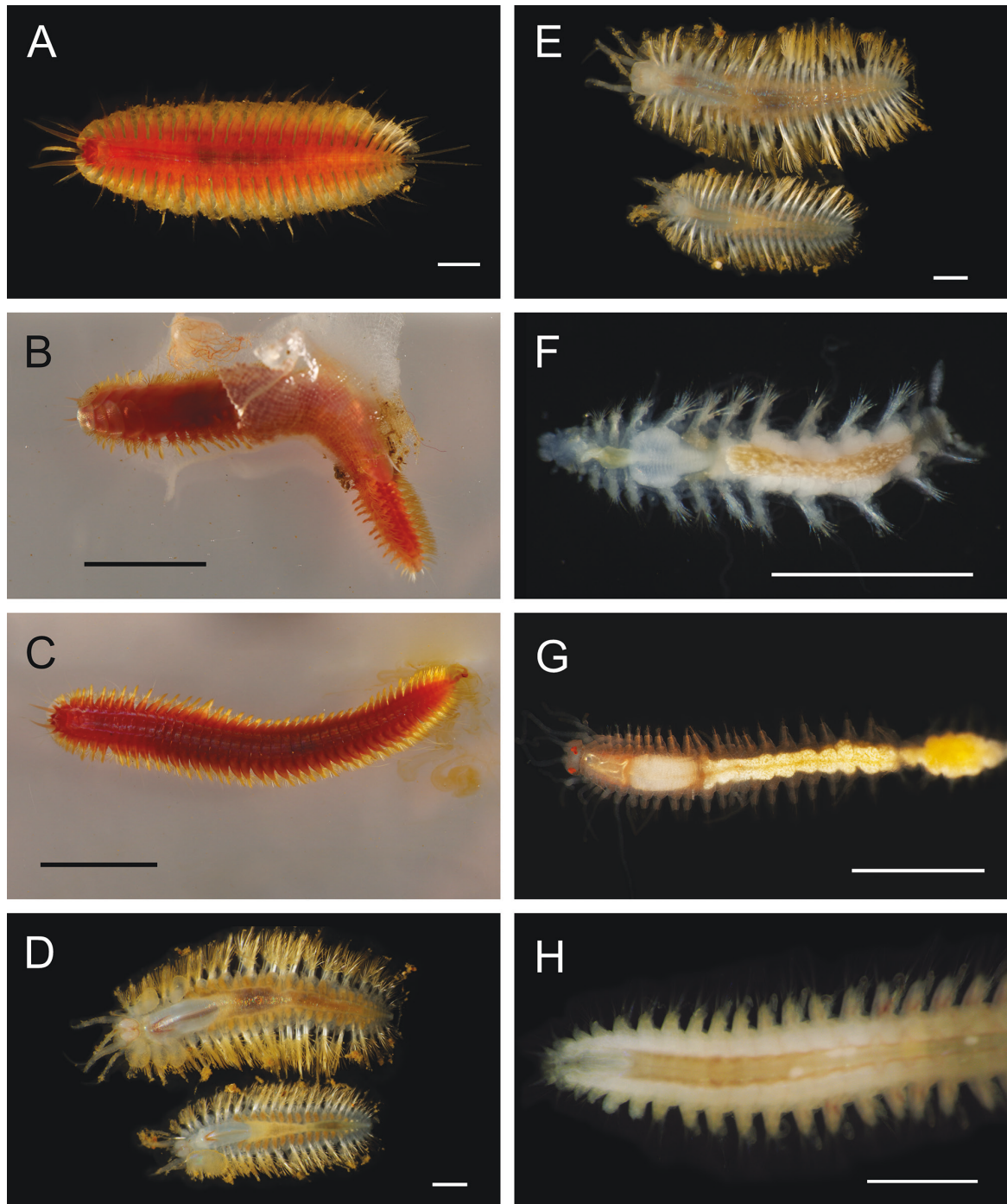


Figure 8. Annelida: Polynoidae, Syllidae, and Nephtyidae, representative live images **A** Polynoidae sp. SIO_BIC_A10082 (A10082, ventral view) **B** Polynoidae sp. SIO_BIC_A10096 (A10096, dorsal view) **C** Polynoidae sp. SIO_BIC_A10096 (A10096, ventral view) **D** Polynoidae sp. SIO_BIC_A10189 (A10189, dorsal view) **E** Polynoidae sp. SIO_BIC_A10189 (A10189, ventral view) **F** *Anguillosyllis* sp. SIO_BIC_A12403 (A12403) **G** *Synmerosyllis* stet. (A1928) **H** *Nephtys* stet. (A1437, dorsal view). Scale bars: 1 mm (**A**, **D–H**); 1 cm (**B**, **C**).

***Synmerosyllis* stet.**

Fig. 8G

Reference. Aguado et al. (2012), in which the published DNA sequences ([JF903573](#), [JF903681](#), [JF903759](#)) correspond to specimen A1928 from AD4588.

Localities. Mound 12 (997 m).

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida |
Nephtyiformia | Nephtyidae

Nephtys stet.

Figs 8H, 9A

Material examined. AD4510: A1437.

Localities. Jacó Summit (742 m).

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida |
Nephtyiformia | Pilargidae

***Sigambra* sp. SIO_BIC_A9597**

Fig. 9B

Material examined. AD4503: A1346; AD4972: A9597; AD4987: A9843.

Localities. Mound 12 (~ 967–999 m), Jacó Scar (1795 m).

Remarks. An undescribed species.

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida | Hesionoidea
| Chrysopetalidae

***Chrysopetalinae* sp. SIO_BIC_A8064**

Fig. 9C

Material examined. AD4508: A1404, A2410; AD4914: A8063; AD4917: A8065;
AD4922: A8064, A10280; AD4974: A9609, A9758; AD4985: A9853; AD4990:
A9879.

Localities. Mound 12 (~ 965–1002 m), Parrita Seep (1401–1402 m), Jacó
Scar (~ 1632–1886 m).

Remarks. An undescribed genus and species.

***Laubierus alvini* Aguado & Rouse, 2011**

Fig. 9D

Reference. Aguado and Rouse 2011**.

Additional material examined. S0214: A10057 ([PQ449257](#)).

Localities. Mound 12 (~ 982–999 m), Mound 11 (~ 1004–1011 m; type lo-
cality), Jacó Scar (~ 1752–1860 m).

Distribution. Known only from the CRM seeps.

Remarks. As noted in the original description, *L. alvini* is a symbiont of
the mussel *Bathymodiolus earlougheri*, but not of the two sympatric species
subsequently described as *B. billschneideri* and *B. nancyschneiderae*. The
worms are found among the gill filaments, with typically 2–25 individuals per
host, and the number of individuals linearly correlates with the length of the
host (Aguado and Rouse 2011).

***Micospina auribohnorum* Watson, Carvajal, Sergeeva, Pleijel & Rouse, 2016**

Fig. 9E

Reference. Watson et al. 2016**.

Additional material examined. S0217: A10088 ([PQ449265](#)).

Localities. Jacó Summit (~ 750 m), Mound 12 (~ 1000 m), Mound 11 (~ 1040 m), The Thumb (1072 m; this study), Jacó Scar (~ 1800 m).

Distribution. Also known from a whale fall at 845 m off San Diego, California (type locality) (Watson et al. 2016).

***Natsushima sashai* Aguado & Rouse, 2011**

Fig. 9F

Reference. Aguado and Rouse 2011**.

Localities. Mound 12 (~ 1000 m; type locality).

Distribution. Known only from the CRM seeps.

Remarks. A symbiont of the solemyid clam *Acharax* cf. *johnsoni*; the worms are found among the gill lamellae, with no more than four individuals per host, in three possible combinations: one female, one male and one female, or two males and two females (Aguado and Rouse 2011).

***Shinkai fontefridae* Aguado & Rouse, 2011**

Fig. 9G

Reference. Aguado and Rouse 2011**.

Localities. Parrita Seep (1186 m), Parrita Scar (~ 1660 m), Jacó Scar (~ 1752–1800 m; type locality).

Distribution. Known only from the CRM seeps.

Remarks. Symbiont of the vesicomylid clams *Phreagena soyoae* and *Archivesica gigas* (previously cited as *Calyptogena kilmeri* and *Vesicomya gigas*, respectively); the worms are found between the gill lamellae and foot, typically with one male and one female per host (Aguado and Rouse 2011).

***Shinkai longipedata* Miura & Ohta, 1991**

Fig. 9H

Reference. Aguado and Rouse 2011.

Localities. Mound 11 (~ 1019–1025 m).

Distribution. Originally described from hydrothermal vents at Iheya Ridge off southern Japan (Miura and Ohta 1991). The occurrences of *S. longipedata* at the CRM established the first record of this species in the eastern Pacific, based on morphological similarity to specimens from Japan and in the absence of DNA sequences from the type locality (Aguado and Rouse 2011). Genetic data from the type locality will be necessary to confirm the apparent trans-Pacific distribution of this species at both vents and seeps.

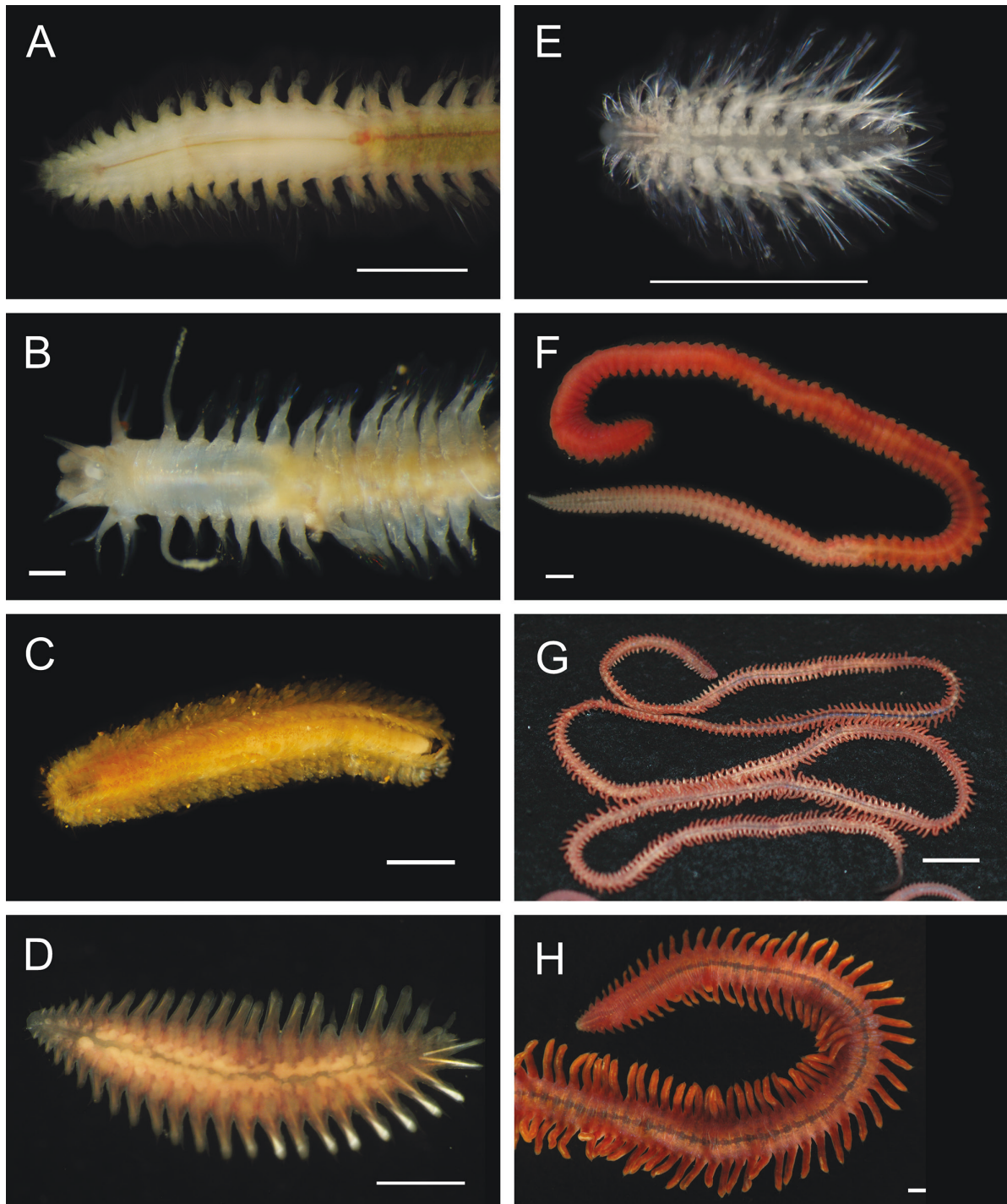


Figure 9. Annelida: Nephtyidae, Pilargidae, and Chrysopetalidae, representative live images **A** *Nephtys stet.* (A1437, ventral view) **B** *Sigambra* sp. SIO_BIC_A9597 (A9597) **C** Chrysopetalinae sp. SIO_BIC_A8064 **D** *Laubierus alvini* (A1323) **E** *Micospina auribohorum* (A1427) **F** *Natsushima sashai* (A1447) **G** *Shinkai fontefridae* (A1384) **H** *Shinkai longipedata* (A1360). Scale bars: 1 mm (**A–F, H**); 1 cm (**G**).

Remarks. Originally described as a commensal symbiont in the mantle cavity of the vesicomid clam *Calyptogena* sp. (Miura and Ohta 1991). At the CRM seeps, the host is the vesicomid clam *Phreagena soyoae* (previously identified as an “undescribed” vesicomid); the worms are found between the gill lamellae and foot, typically with one male and one female per host (Aguado and Rouse 2011).

***Vigtorniella* sp. SIO_BIC_A8061**

Fig. 10A

Material examined. AD4914: A8061, A8062.

Localities. Jacó Scar (~ 1632–1886 m).

Remarks. An undescribed species.

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida | Hesionoidea
| Hesionidae

***Amphiduropsis* cf. *axialensis* (Blake & Hilbig, 1990)**

Fig. 10B

Reference. Rouse et al. 2018.

Localities. Mound 12 (~ 984–997 m).

Remarks. The type locality of *Amphiduropsis axialensis* is hydrothermal vents at 1545 m on the Axial Seamount of the Juan de Fuca Ridge (Blake and Hilbig 1990). As discussed in Rouse et al. (2018), the specimens above (and other seep specimens from Hydrate Ridge off Oregon, 587–809 m, and the Guaymas Basin, Gulf of California, 1560–1613 m) are morphologically consistent with *A. axialensis* but are conservatively designated as *A. cf. axialensis* because DNA sequences are not available from the type locality.

***Gyptis robertscripsi* Rouse, Carvajal & Pleijel, 2018**

Fig. 10C

Reference. Rouse et al. 2018**.

Localities. Mound 12 (~ 982–1002 m; type locality), Mound 11 (~ 1019–1025 m), Jacó Scar (1783–1794 m).

Distribution. Known only from the CRM seeps.

***Gyptis* sp. SIO_BIC_A10083**

Fig. 10D

Material examined. AD4919: A8114; S0216: A10083.

Localities. Quepos Slide (~ 379–398 m).

Remarks. An undescribed species.

***Leocrates* gen. inc.**

Fig. 10E

Material examined. AD4914: A8113; S0230: A10184.

Localities. Jacó Scar (1886 m), Mound Jaguar (1909 m).

Remarks. Taxonomic placement of these specimens requires further assessment, particularly in light of recent revisions to several hesionid genera (Salazar-Vallejo 2020b).

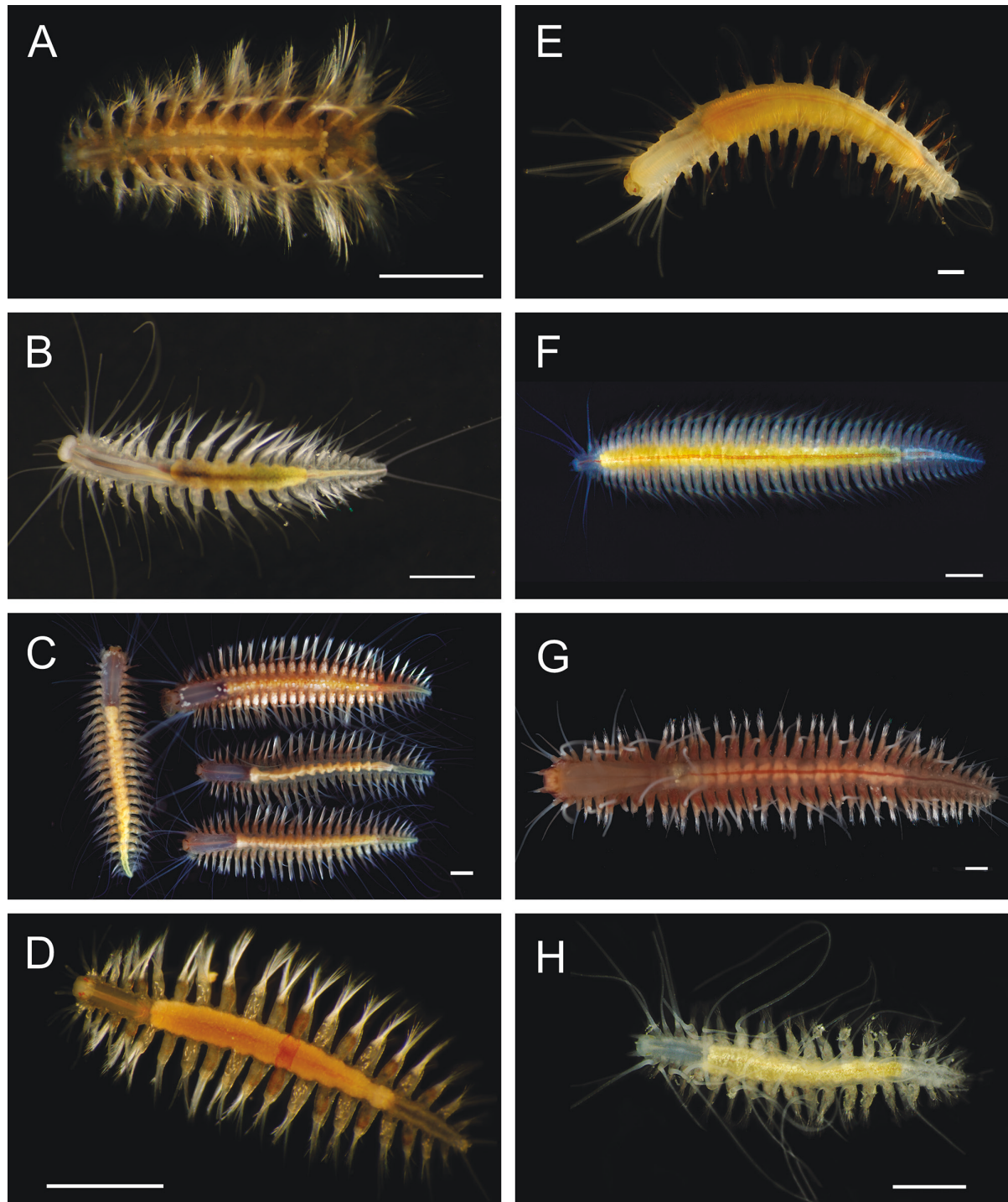


Figure 10. Annelida: Chrysopetalidae and Hesionidae, representative live images **A** *Vigtorniella* sp. SIO_BIC_A8061 (A8061) **B** *Amphiduopsis* cf. *axialensis* (A8110) **C** *Gyptis robertscrippsi* (A1750) **D** *Gyptis* sp. SIO_BIC_A10083 (A10083) **E** *Leocrates* gen. inc. (A10184) **F** *Neogyptis jeffruoccoi* (A1448) **G** *Sirsoe dalailamai* (MZUCR 401-01) **H** *Sirsoe munki* (A1409). Scale bars: 1 mm.

***Neogyptis jeffruoccoi* Rouse, Carvajal & Pleijel, 2018**

Fig. 10F

Reference. Rouse et al. 2018**.

Localities. Mound 12 (~ 988–997 m; type locality), Mound 11 (~ 1019–1025 m).

Distribution. Also known from seeps in the Guaymas Basin, Gulf of California, 1572–1613 m, and off Del Mar, California, 1020 m (Rouse et al. 2018).

Remarks. Found in the mantle cavity of the solemyid clam *Acharax* cf. *johnsoni*, with either one individual or two (one male and one female) per host (Rouse et al. 2018).

***Sirsoe dalailamai* Rouse, Carvajal & Pleijel, 2018**

Fig. 10G

Reference. Rouse et al. 2018**.

Localities. Mound 12 (997 m), Parrita Seep (1402 m), Jacó Scar (1784–1795 m; type locality).

Distribution. Also described from a seep in the Guaymas Basin, Gulf of California, 1560–1613 m (Rouse et al. 2018).

Remarks. Associated with vestimentiferan and mussel communities at areas of active methane seepage (Rouse et al. 2018).

***Sirsoe munki* Rouse, Carvajal & Pleijel, 2018**

Fig. 10H

Reference. Rouse et al. 2018**.

Additional material examined. S0217: A10087 ([PQ449264](#)).

Localities. Jacó Slope (1063 m), The Thumb (1072 m; this study), Jacó Scar (~ 1800 m; type locality).

Distribution. Known only from the CRM seeps.

***Sirsoe* sp. SIO_BIC_A8288**

Fig. 11A

Material examined. AD4909: A8288 ([PQ449293](#)).

Localities. Mound 12 (990 m).

Remarks. An undescribed species associated with a microbial mat.

***Vrijenhoekia* sp. A sec. Summers et al. 2015**

Fig. 11B

Material examined. AD4508: A1406; AD4907: A8101; AD4974: A9606, A9608, A9612, A9615; AD4988: A9926.

Localities. Mound 11 (1010 m), Mound 12 (~ 990–999 m), Parrita Seep (1419 m).

Remarks. The undescribed species *Vrijenhoekia* sp. A was previously known only from a single specimen, SIO-BIC A3255, from a seep in the Guaymas Basin, Gulf of California, 1565–1598 m; the specimen was collected in poor condition and diagnostic morphological features could not be assessed (Summers et al. 2015). The CRM specimens represent the first records of this morphospecies beyond the Guaymas Basin and expand the known depth range to ~ 1000–1600 m. The CRM specimens were associated with naturally occurring and experimentally deployed wood (A1406, A9606, A9608, A9926) or experimentally deployed bone (A9615).

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida | Nereididae

These specimens will be further examined and compared to other eastern Pacific Nereididae in a separate work.

***Nereis* stet.**

Fig. 11C

Material examined. AD4906: A8258; AD4910: A8291.

Localities. Mound 12 (997–1002 m).

Remarks. A8258 was associated with experimentally deployed bone.

Nereididae sp. SIO_BIC_A9614

Fig. 11D

Material examined. AD4974: A9614.

Localities. Mound 12 (992 m)

Remarks. An undescribed species. Associated with experimentally deployed wood and pig bones.

***Pectinereis strickrotti* Villalobos-Guerrero, Huč, Tilic, Hiley & Rouse, 2024**

Fig. 11E

Reference. Villalobos-Guerrero et al. 2024**.

Material examined. AD4984: A9889.

Localities. Mound 12 (996–1010 m).

Distribution. Known only from the CRM seeps.

Remarks. Known from epitokous males, all collected while swimming just above the seafloor, and from a fragment of an atokous infaunal female recovered from a push core sample (Villalobos-Guerrero et al. 2024).

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida |
Glyceriformia | Sphaerodoridae

***Sphaerephesia* sp. SIO_BIC_A10069**

Fig. 11F

Material examined. S0214: A10069 ([PQ449261](#); no voucher remaining).

Localities. Jacó Scar (1875 m).

Remarks. Associated with a naturally occurring wood fall. The closest COI BLASTN result on GenBank was a specimen of *Sphaerephesia* cf. *discolis* (Borowski, 1994) from the Brazilian Basin, 5180 m ([KR019875.1](#), 95.39% identity, formerly *Sphaerodoropsis* cf. *discolis*) (Capa et al. 2016, 2019). *Sphaerephesia discolis* was originally described from the Peru Basin, 4152 m (Capa et al. 2019), so a comparison of the CRM specimen to material from the type locality would be informative.

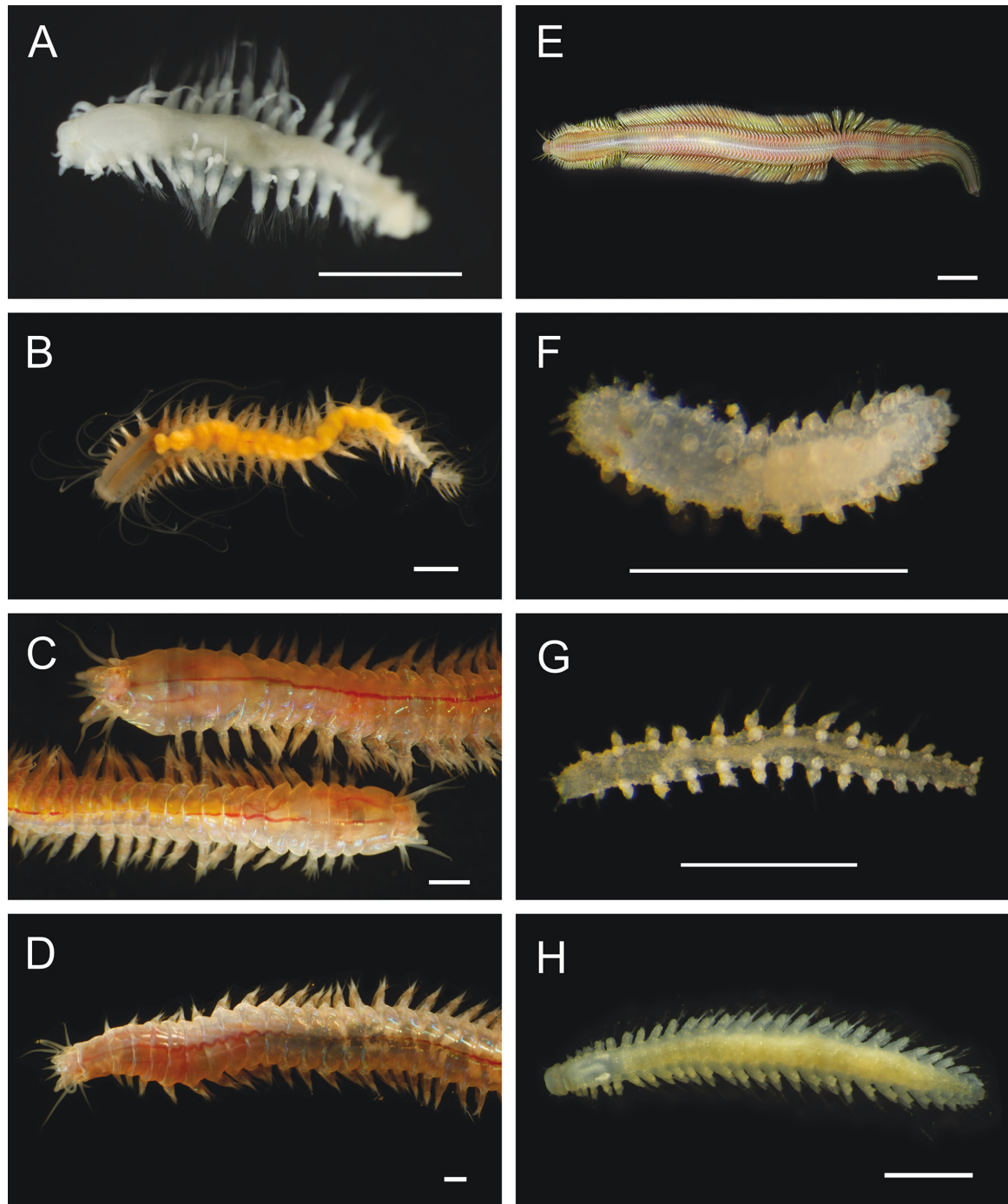


Figure 11. Annelida: Hesionidae, Nereididae, Sphaerodoridae, and Lacydoniidae, representative images. Live specimens are depicted unless otherwise specified **A** *Sirsoe* sp. SIO_BIC_A8288 (A8288, preserved specimen) **B** *Vrijenhoekia* sp. A sec. Summers et al. 2015 (A9606) **C** *Nereis* stet. (A8291) **D** Nereididae sp. SIO_BIC_A9614 (A9614) **E** *Pectinereis strickrotti* (A9889) **F** *Sphaerephesia* sp. SIO_BIC_A10069 (A10069) **G** *Sphaerodoropsis* sp. SIO_BIC_A10068 (A10068) **H** *Lacydonia* sp. SIO_BIC_A1432 (A1432). Scale bars: 1 mm (**A–D, F–H**); 1 cm (**E**).

***Sphaerodoropsis* sp. SIO_BIC_A10068**

Fig. 11G

Material examined. S0214: A10068 ([PQ449260](#)).

Localities. Jacó Scar (1875 m).

Remarks. Associated with a naturally occurring wood fall.

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida |
Phyllodociformia

These specimens will be further examined and compared to other eastern Pacific *Lacydonia* in a separate work.

***Lacydonia* sp. SIO_BIC_A1432**

Fig. 11H

Material examined. AD4504: A1525, A1526; AD4510: A1432.

Localities. Jacó Summit (~ 741–744 m), Mound 11 (~ 1004–1011 m).

Remarks. An undescribed species.

***Lacydonia* sp. SIO_BIC_A1606**

Material examined. AD4510: A1606.

Localities. Jacó Summit (~ 741–744 m).

Remarks. An undescribed species.

***Lacydonia* sp. SIO_BIC_A9774**

Fig. 12A

Material examined. AD4588: A1925; AD4978: A9774; AD4990: A9880; S0217: A10090 ([MZ562520](#)).

Localities. The Thumb (~ 940–1070 m), Mound 12 (~ 996–999 m), Parrita Seep (1401 m).

Remarks. An undescribed species.

***Lacydonia* sp. SIO_BIC_A16347**

Material examined. AD4504: A16347.

Localities. Mound 11 (~ 1004–1011 m).

Remarks. An undescribed species.

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida |
Phyllodociformia | Phyllodocidae

***Eulalia* sp. SIO_BIC_A1383**

Fig. 12B, C

Material examined. AD4506: A1383 ([PQ449278](#)).

Localities. Parrita Seep (~ 1030–1179 m).

***Galapagomystides patricki* Pearson & Rouse, 2022**

Fig. 12D

Reference. Pearson and Rouse 2022**.

Localities. Parrita Seep (~ 1401–1419 m; type locality), Jacó Scar (1762–1785 m).

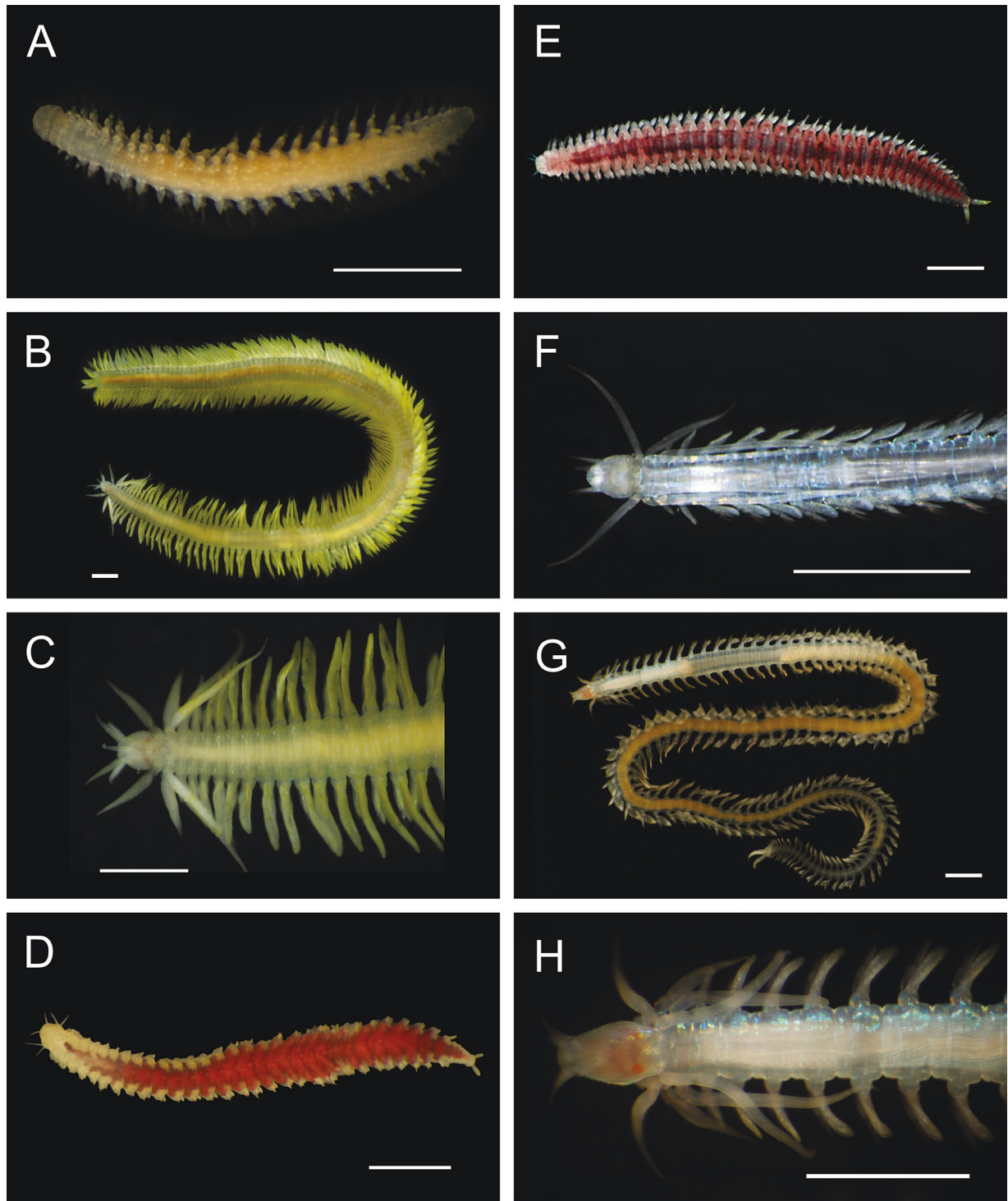


Figure 12. Annelida: Lacydoniidae and Phyllodoceidae, representative live images **A** *Lacydonia* sp. SIO_BIC_A9774 (A9774) **B** *Eulalia* sp. SIO_BIC_A1383 (A1383, wide view) **C** *Eulalia* sp. SIO_BIC_A1383 (A1383, anterior detail) **D** *Galapagomystides patricki* (A9934) **E** *Galapagomystides verenae* (A10044) **F** *Phyllodoce* sp. SIO_BIC_A1469 (A1469) **G** *Phyllodoce* sp. SIO_BIC_A8454 (A8454, wide view) **H** *Phyllodoce* sp. SIO_BIC_A8454 (A8454, anterior detail). Scale bars: 1 mm.

Distribution. Known only from the CRM seeps and vicinity. One paratype was collected from a multicore sample ca 21 km northwest of Parrita Seep, at 805 m depth (Cruise AT15-44, Multicore 17: 9.1713, -84.7459) (Pearson and Rouse 2022).

Remarks. Some specimens were associated with empty vestimentiferan tubes (Pearson and Rouse 2022).

***Galapagomystides verenae* (Blake & Hilbig, 1990)**

Fig. 12E

Reference. Pearson and Rouse 2022.

Localities. Mound 12 (~ 984–997 m), Parrita Seep (~ 1400–1410 m), Jacó Scar (~ 1632–1908 m).

Distribution. Originally described from vents on the Juan de Fuca and Explorer Ridges, 1545–2195 m (Blake and Hilbig 1990). Also known from seeps in the Guaymas Basin, Gulf of California, to the CRM, 1572 to ~ 1800 m (Pearson and Rouse 2022).

Remarks. At the CRM seeps, *G. verenae* is associated with the tubes of juvenile *Escarpia spicata* and may feed on the blood of vestimentiferans (Pearson and Rouse 2022).

***Phyllodoce* sp. SIO_BIC_A1469**

Fig. 12F

Material examined. AT15-44 MC1-2: A1469 ([PQ449283](#)).

Localities. Near Mound 12 (1019 m).

Remarks. This specimen was collected in a sediment core adjacent to Mound 12, ca 500 m from known sites of active seepage and likely representing the far-transition zone to the surrounding environment.

***Phyllodoce* sp. SIO_BIC_A8454**

Fig. 12G, H

Material examined. AD4505: A1366; AD4508: A1399; AD4922: A8454 ([PQ449312](#)); AD4975: A9795; AD4978: A9798 ([PQ449326](#)).

Localities. Mound 12 (~ 964–1009 m), Mound 11 (~ 1019–1025 m), Parrita Seep (~ 1401–1419 m).

Phyllodocidae sp. SIO_BIC_A10054

Fig. 13A

Material examined. S0212: A10054 ([PQ449256](#)).

Localities. Jacó Scar (~ 1780–1860 m).

***Sige* sp. SIO_BIC_A8263**

Fig. 13B, C

Material examined. AD4906: A8263 ([PQ449290](#)); AD4912: A8356 ([PQ449301](#)).

Localities. Mound 12 (1002 m), Jacó Scar (~ 1795–1859 m).

***Sige* sp. SIO_BIC_A8430**

Fig. 13D

Material examined. AD4907: A8284 ([PQ449292](#)); AD4922: A8430 ([PQ449311](#)).

Localities. Mound 12 (~ 990–1002 m).

Remarks. A8430 was associated with a naturally occurring wood fall.

Annelida | Polychaeta | Errantia | Aciculata | Phyllodocida |
Phyllodociformia

***Paralacydonia* stet.**

Fig. 13E

Material examined. AD4509: A1412.

Localities. Jacó Scar (1855 m).

Annelida | Polychaeta | Errantia | Aciculata | Eunicida | Oenonoidea

Lumbrineridae stet.

Fig. 13F

Material examined. AD4505: A1370 ([PQ448999](#)).

Localities. Mound 11 (~ 1019–1025 m).

Annelida | Polychaeta | Errantia | Aciculata | Eunicida | Eunicoidea |
Eunicidae

Eunicidae stet.

Fig. 13G

Material examined. AD4510: A1431; AD4914: A8386; AD4985: A9895
([PQ449332](#)).

Localities. Mound 12 (991 m), Jacó Summit (741 m), Jacó Scar (1798 m).

Remarks. The closest COI BLASTN result on GenBank was *Leodice antarctica* (Baird, 1869) ([GQ497532.1](#), 92.64% identity, formerly *Eunice antarctica*), known from Antarctic and sub-Antarctic waters (Baird 1869). The identification of species of *Eunice* and *Leodice*, among other genera, is complicated by the need for taxonomic revision (Zanol et al. 2021).

Annelida | Polychaeta | Errantia | Aciculata | Eunicida | Eunicoidea |
Onuphidae

***Hyalinoecia* stet.**

Fig. 13H

Material examined. AD4510: A1440; S0214: A10060 ([PQ449259](#)).

Localities. Jacó Summit (~ 741–744 m), Jacó Scar (1875 m).

Annelida | Polychaeta | Errantia | Aciculata | Eunicida | Dorvilleidae

***Ophryotrocha* cf. *batillus* Wiklund et al., 2012**

Fig. 14A

Material examined. AD4974: A9610 (MT444001).

Localities. Mound 12 (992 m).

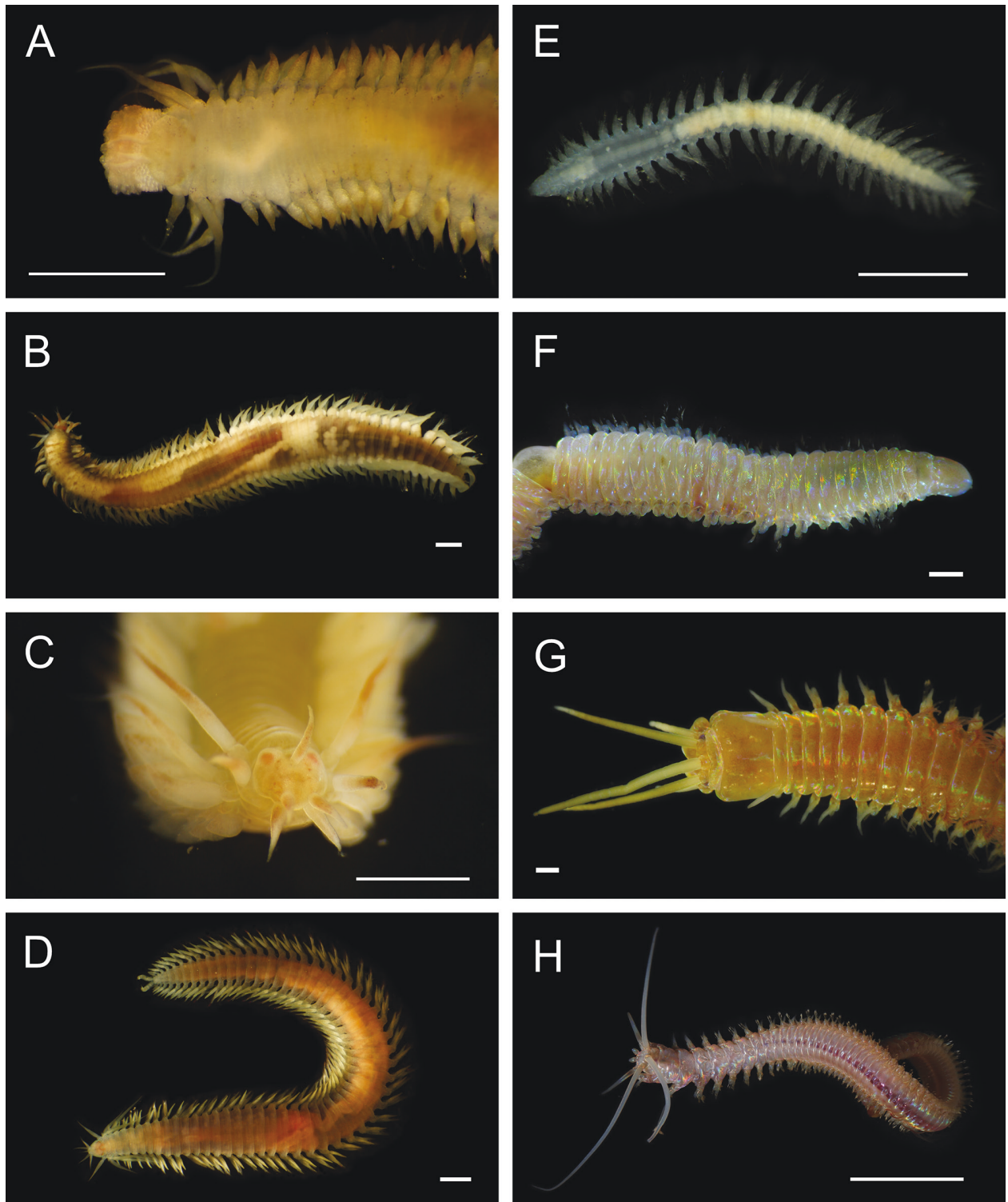


Figure 13. Annelida: Phyllodocidae, *Paralacydonia*, Eunicidae, and Onuphidae, representative live images **A** Phyllodocidae sp. SIO_BIC_A10054 (A10054) **B** *Sige* sp. SIO_BIC_A8263 (A8263) **C** *Sige* sp. SIO_BIC_A8263 (A8356) **D** *Sige* sp. SIO_BIC_A8430 (A8430) **E** *Paralacydonia* stet. (A1412, wide view) **F** Lumbrineridae stet. (A1370) **G** Eunicidae stet. (A1431) **H** *Hyalinoecia* stet. (A10060). Scale bars: 1 mm (**A–G**); 1 cm (**H**)

Remarks. Associated with experimentally deployed wood. To be discussed in a forthcoming study. *Ophryotrocha batillus* was originally described from whale falls and wood falls off southern California, 960–1960 m (Wiklund et al. 2012).

***Ophryotrocha* cf. *flabella* Wiklund et al., 2012**

Fig. 14B

Material examined. AD4509: A1410 ([MT435616](#)); AD4988: A9928 ([MT435618](#); no specimen remaining), A9929 ([MT435617](#)).

Localities. Mound 11 (1010 m), Jacó Slope (1064 m).

Remarks. To be discussed in a forthcoming study. *Ophryotrocha flabella* was originally described from whale falls off southern California, 960–1960 m (Wiklund et al. 2012). Specimens A9928 and A9929 were associated with a naturally occurring wood fall.

***Ophryotrocha* cf. *platycephale* Blake, 1985**

Fig. 14C

Material examined. AD4990: A9878 ([MT435620](#)).

Localities. Parrita Seep (~ 1400–1435 m).

Remarks. To be discussed in a forthcoming study. *Ophryotrocha platycephale* was originally described from sedimented vents at 2000–2030 m in the Guaymas Basin, Gulf of California (Blake 1985; Solís-Weiss and Hilbig 1992).

***Ophryotrocha* sp. SIO_BIC_A8367**

Fig. 14D

Material examined. AD4914: A8367 ([PQ449305](#)).

Localities. Jacó Scar (~ 1632–1886 m).

Remarks. Likely an undescribed species.

***Ophryotrocha* sp. SIO_BIC_A9611**

Fig. 14E

Material examined. AD4974: A9611 ([PQ449317](#)).

Localities. Mound 12 (992 m).

Remarks. An undescribed species, associated with experimentally deployed bone and wood.

***Ophryotrocha* sp. SIO_BIC_A9723**

Fig. 14F

Material examined. AD4976: A9723 ([PQ449322](#)).

Localities. Jacó Scar (1887 m).

Remarks. Likely an undescribed species. Associated with experimentally deployed wood.

***Ophryotrocha* sp. SIO_BIC_A9800**

Fig. 14G

Material examined. AD4912: A8354; AD4913: A8360; AD4978: A9800 ([MT435612](#)).

Localities. Mound 12 (999 m), Jacó Scar (~ 1795–1859 m).

Remarks. An undescribed species.

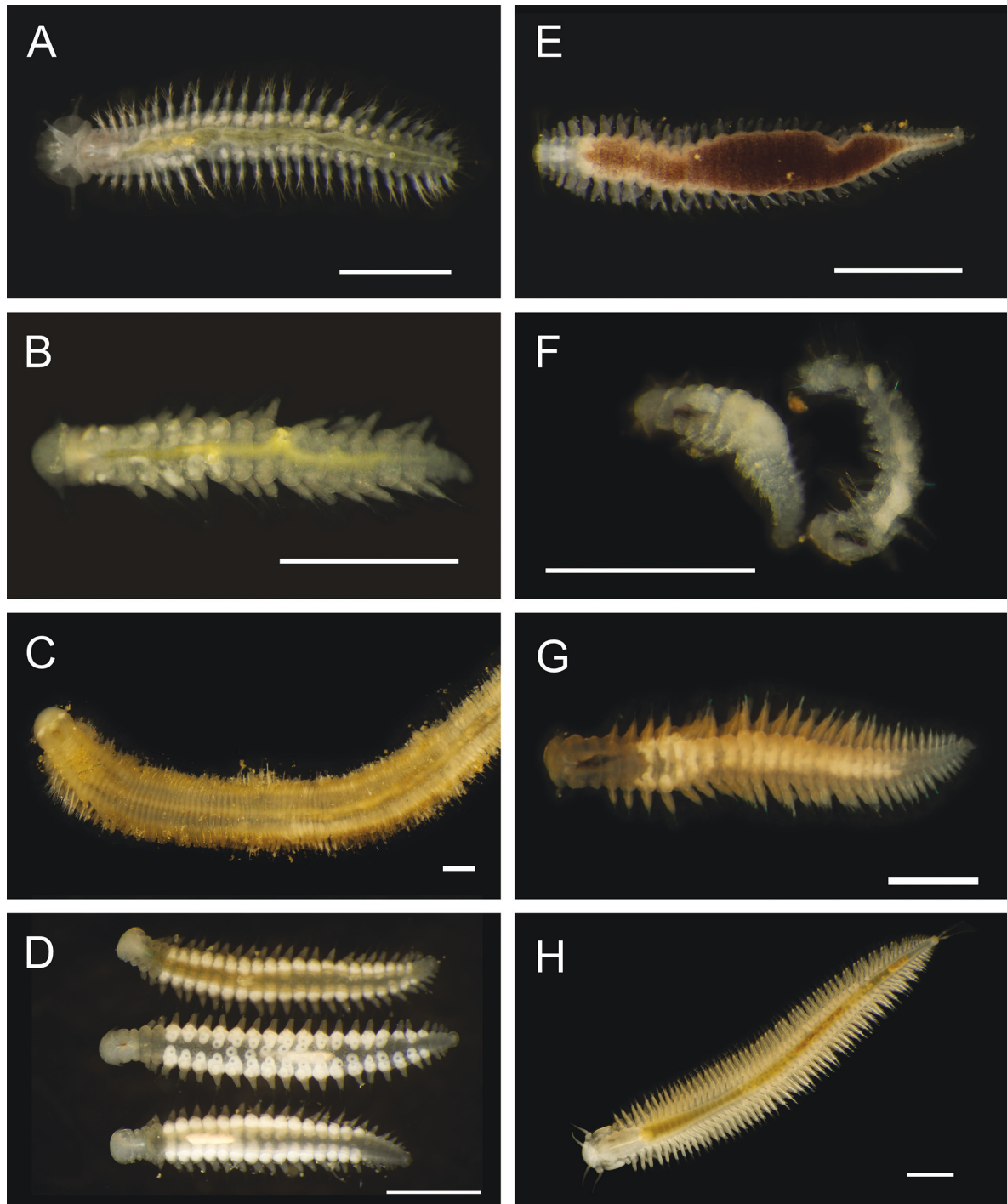


Figure 14. Annelida: Dorvilleidae: *Ophryotrocha*, representative live images **A** *Ophryotrocha* cf. *batillus* (A9610) **B** *Ophryotrocha* cf. *flabella* (A1410) **C** *Ophryotrocha* cf. *platycephale* (A9878) **D** *Ophryotrocha* sp. SIO_BIC_A8367 (A8367) **E** *Ophryotrocha* sp. SIO_BIC_A9611 (A9611) **F** *Ophryotrocha* sp. SIO_BIC_A9723 (A9723) **G** *Ophryotrocha* sp. SIO_BIC_A9800 (A9800) **H** *Ophryotrocha* sp. SIO_BIC_A10052 (A10052). Scale bars: 1 mm.

***Ophryotrocha* sp. SIO_BIC_A10052**

Fig. 14H

Material examined. S0213: A10052 ([MT435579](https://doi.org/10.3897/zookeys.1222.134385)).

Localities. Jacó Summit (~ 730–820 m).

Remarks. An undescribed species.

***Ophryotrocha* sp. SIO_BIC_A10084**

Fig. 15A

Material examined. S0217: A10084 ([MT435566](#)), MZUCR 1512-01; S0219: A10113 ([MT435562](#)).

Localities. Rio Bongo Scar (610 m), The Thumb (1072 m).

Remarks. An undescribed species.

***Ophryotrocha* sp. SIO_BIC_A10106**

Fig. 15B

Material examined. S0219: A10106 ([PQ449268](#)).

Localities. Rio Bongo Scar (661 m).

Remarks. An undescribed species associated with a naturally occurring wood fall.

***Ophryotrocha* sp. SIO_BIC_A10114**

Fig. 15C

Material examined. S0219: A10114 ([MT435596](#)), MZUCR 1511-01.

Localities. Rio Bongo Scar (~ 480–650 m).

Remarks. An undescribed species to be described in a forthcoming study.

***Parougia ceruleibohnorum* Yen & Rouse, 2020**

Fig. 15D

Reference. Yen and Rouse 2020**.

Localities. Mound 12 (996 m), Parrita Seep (~ 1401–1419 m; type locality).

Distribution. Known only from the CRM seeps.

***Parougia* cf. *billiemi*roae Yen & Rouse, 2020**

Fig. 15E

Reference. Yen and Rouse 2020.

Localities. Jacó Scar (1796 m).

Remarks. *Parougia billiemi*roae *sensu stricto* is found at seeps, 587–1583 m, from Hydrate Ridge off Oregon (type locality) to the Guaymas Basin, Gulf of California (Yen and Rouse 2020). As discussed in Yen and Rouse (2020), the CRM single specimen A9678 is morphologically indistinguishable from *P. billiemi*roae *sensu stricto*, but its COI haplotype is up to 5.8% divergent from the type series and has been conservatively designated *P. cf. billiemi*roae.

***Parougia* cf. *sulleyi* Yen & Rouse, 2020**

Fig. 15F

Reference. Yen and Rouse 2020.

Localities. Mound 12 (~ 987–997 m).

Remarks. *P. sulleyi sensu stricto* is found at seeps, 600–1600 m, from Hydrate Ridge off Oregon to the Guaymas Basin, Gulf of California (type locality) (Yen

and Rouse 2020). As discussed in Yen and Rouse (2020), the CRM specimens are morphologically indistinguishable from *P. sulleyi sensu stricto* but share a COI haplotype that is 7.4% divergent from the type series. *Parougia* cf. *sulleyi* is the same taxon as the “*Dorvillea* sp.” documented by Thurber et al. (2012) as a consumer of archaea at the CRM seeps, representing one of the first records of “archivory” by a metazoan (Yen and Rouse 2020).

***Parougia theloniousblueski* Yen & Rouse, 2020**

Fig. 15G

Reference. Yen and Rouse 2020**.

Localities. Mound 12 (987–997 m), Mound 11 (1010 m; type locality).

Distribution. Known only from the CRM seeps.

Annelida | Polychaeta | Errantia | Aciculata incertae sedis |
Myzostomida

***Eenymeenymyzostoma* sp. SIO_BIC_A8428**

Figs 15H, 16A

Material examined. AD4501: A1476; AD4503: A1340; AD4922: A8427 ([PQ449309](#)), A8428, A8431.

Localities. Mound 12 (~ 966–995 m).

Remarks. An undescribed species associated with the antipatharian coral *Lillipathes ritamariae*.

***Myzostoma josefinae* Summers & Rouse in Summers et al. 2014**

Fig. 16B

Material examined. AD4913: A8362.

Localities. Jacó Scar (1878 m).

Distribution. Originally described from a whale fall at 1020 m in Monterey Submarine Canyon, off California (type locality) and in the vicinity of sedimented vents and seeps at ~ 1350 m in the Guaymas Basin, Gulf of California (Summers et al. 2014b).

New records. The CRM specimen represents a new southern record and a new maximum depth record for this species.

Remarks. The specimen, showing the distinctive paired elongate caudal appendages, was associated with the crinoid *Psathyrometra* cf. *fragilis* (E7034), consistent with previous records of *M. josefinae* on *P. fragilis* (Summers et al. 2014b).

***Pulvinomyzostomum inaki* Summers & Rouse in Summers et al. 2014**

Fig. 16C

Reference. Summers et al. 2014b**.

Localities. Jacó Scar (1789 m; type locality).

Distribution. Known only from the CRM seeps.

Remarks. Associated with an antedonid crinoid (E4399) (Summers et al. 2014b).

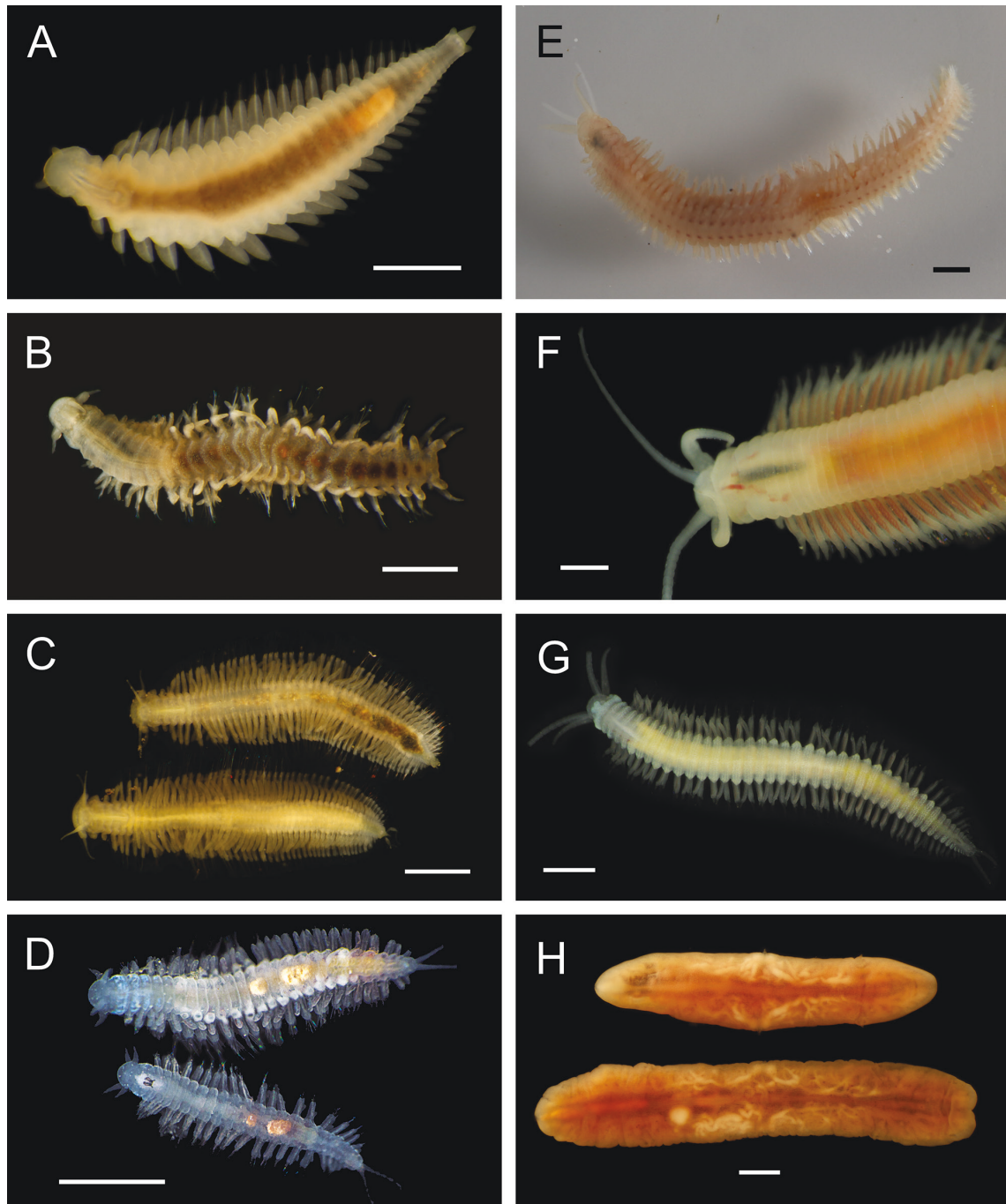


Figure 15. Annelida: Dorvilleidae and Myzostomida, representative live images **A** *Ophryotrocha* sp. SIO_BIC_A10084 (A10084) **B** *Ophryotrocha* sp. SIO_BIC_A10106 (A10106) **C** *Ophryotrocha* sp. SIO_BIC_A10114 (A10114) **D** *Parougia ceruleibohnorum* (A1446) **E** *Parougia* cf. *billiemi* (A9678) **F** *Parougia* cf. *sulleyi* (A1333) **G** *Parougia theloniousblueski* (A1337) **H** *Eenyumyzostoma* sp. SIO_BIC_A8428 (A8428, dorsal view). Scale bars: 1 mm.

***Pulvinomyzostomum* sp. SIO_BIC_A8361**

Fig. 16D, E

Material examined. AD4913: A8361 ([PQ449303](#)).

Localities. Jacó Scar (1878 m)

Remarks. An undescribed species. One female and one male were associated with the crinoid *Psathyrometra* cf. *fragilis* (E7034).

Annelida | Polychaeta | Errantia | Aciculata incertae sedis | Nerillidae

Nerillidae stet.

Fig. 16F

Material examined. AD4979: A9797.

Localities. Quepos Slide (~ 380–395 m).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Sabellida | Sabellidae

***Bispira* sp. SIO_BIC_A9587**

Figs 16G, H, 17A

Reference. Goffredi et al. (2020) for characterization of bacterial symbiosis; specimen A9587 was erroneously cited as A9598.

Material examined. AD4508: A1396; AD4513: A1460 ([PQ449282](#); 18S: [PQ304645](#)); AD4916: A8387; AD4971: A9587.

Localities. Parrita Seep (1402 m; this study), Jacó Scar (~ 1604–1854 m).

Remarks. An undescribed species, abundant at zones of active seepage and utilizing methanotrophic bacterial symbionts for nutrition (Goffredi et al. 2020).

***Chone* sp. SIO_BIC_A8422**

Fig. 17B

Material examined. AD4918: A8419; AD4919: A8422.

Localities. Quepos Slide (~ 379–410 m).

Remarks. Likely an undescribed species (María Ana Tovar-Hernández, pers. comm. 8 August 2020).

***Chone* sp. SIO_BIC_A8462**

Fig. 17C

Material examined. AD4916: A8389; AD4919: A8462 (16S: [PQ304651](#)).

Localities. Parrita Seep (~ 1400–1410 m), Jacó Scar (1746 m).

Remarks. Likely an undescribed species (María Ana Tovar-Hernández, pers. comm. 8 August 2020).

***Jasmineira* stet.**

Fig. 17D

Material examined. AD4907: A8282.

Localities. Mound 12 (999 m).

***Pseudopotamilla* sp. SIO_BIC_A1455**

Fig. 17E

Material examined. AD4512: A1455; AD4918: A8418; AD4979: A9796.

Localities. Quepos Slide (~ 338–411 m).

Remarks. An undescribed species.

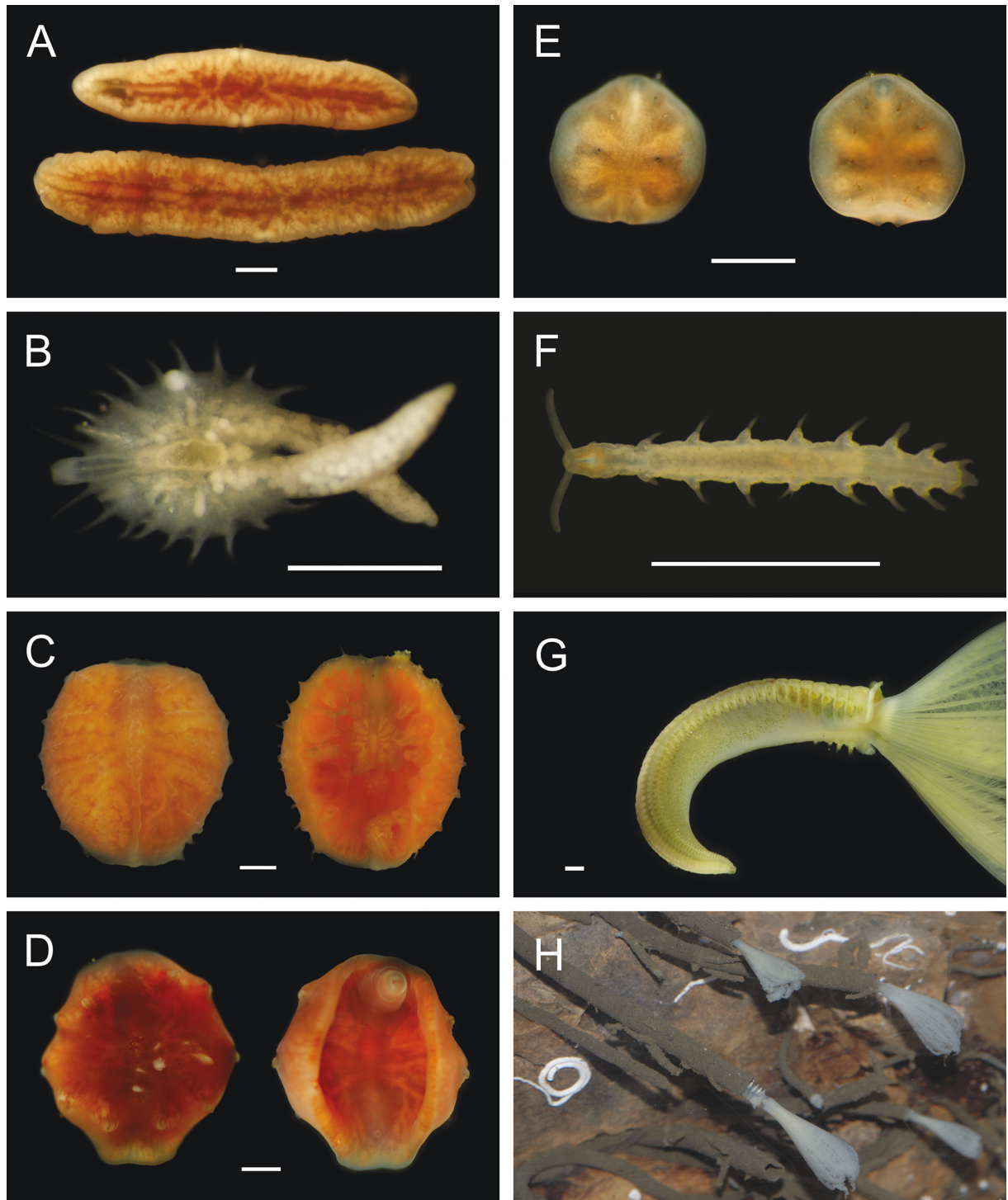


Figure 16. Annelida: Myzostomida, Nerillidae, and Sabellidae, representative live images **A** *Eenymeenymyzostoma* sp. SIO_BIC_A8428 (A8428, ventral view) **B** *Myzostoma josefinae* (A8362) **C** *Pulvinomyzostomum inaki* (A1579, dorsal and ventral views of female with dwarf males) **D** *Pulvinomyzostomum* sp. SIO_BIC_A8361 (A8361, dorsal and ventral views of female) **E** *Pulvinomyzostomum* sp. SIO_BIC_A8361 (A8361, dorsal and ventral views of male) **F** Nerillidae stet. (A9797) **G** *Bispira* sp. SIO_BIC_A9587 (A1396) **H** *Bispira* sp. SIO_BIC_A9587 (A1460, in tubes attached to rocks photographed *ex situ* in the shipboard laboratory). Scale bars: 1 mm.

***Pseudopotamilla* sp. SIO_BIC_A9732**

Fig. 17F

Material examined. AD4978: A9732.

Localities. Mound 12 (~ 996–999 m).

***Sabellidae* sp. SIO_BIC_A8286**

Fig. 17G

Material examined. AD4908: A8286 (18S: [PQ304646](#)), A8287.

Localities. Mound 12 (1000 m).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Sabellida |
Serpulidae

***Hyalopomatus* sp. SIO_BIC_A1434**

Fig. 17H

Reference. Rouse and Kupriyanova (2021) for DNA sequences and phylogenetic analysis.

Material examined. AD4510: A1434.

Localities. Jacó Summit (741–745 m).

Remarks. Possibly an undescribed species (Rouse and Kupriyanova 2021).

***Laminatubus joycebrooksae* Rouse & Kupriyanova, 2021**

Fig. 18A

Reference. Rouse and Kupriyanova 2021**.

Localities. Mound 12 (~ 1000 m; type locality).

Distribution. Known only from the CRM seeps.

***Laminatubus paulbrooksi* Rouse & Kupriyanova, 2021**

Fig. 18B

Reference. Rouse and Kupriyanova 2021**.

Localities. Parrita Seep (1402 m), Jacó Scar (~ 1800 m; type locality).

Distribution. Also known from seeps in the Guaymas Basin and Pescadero Basin, Gulf of California, 1565–2478 m (Rouse and Kupriyanova 2021).

Remarks. Abundant at zones of active seepage, this species utilizes methanotrophic bacterial symbionts for nutrition (Goffredi et al. 2020).

***Protis* stet.**

Material examined. AD4506: A1553 (18S: [PQ304668](#); no image available).

Localities. Parrita Seep (1186 m).

Remarks. We thank Elena Kupriyanova (Australian Museum Research Institute) for generating the 18S sequence.

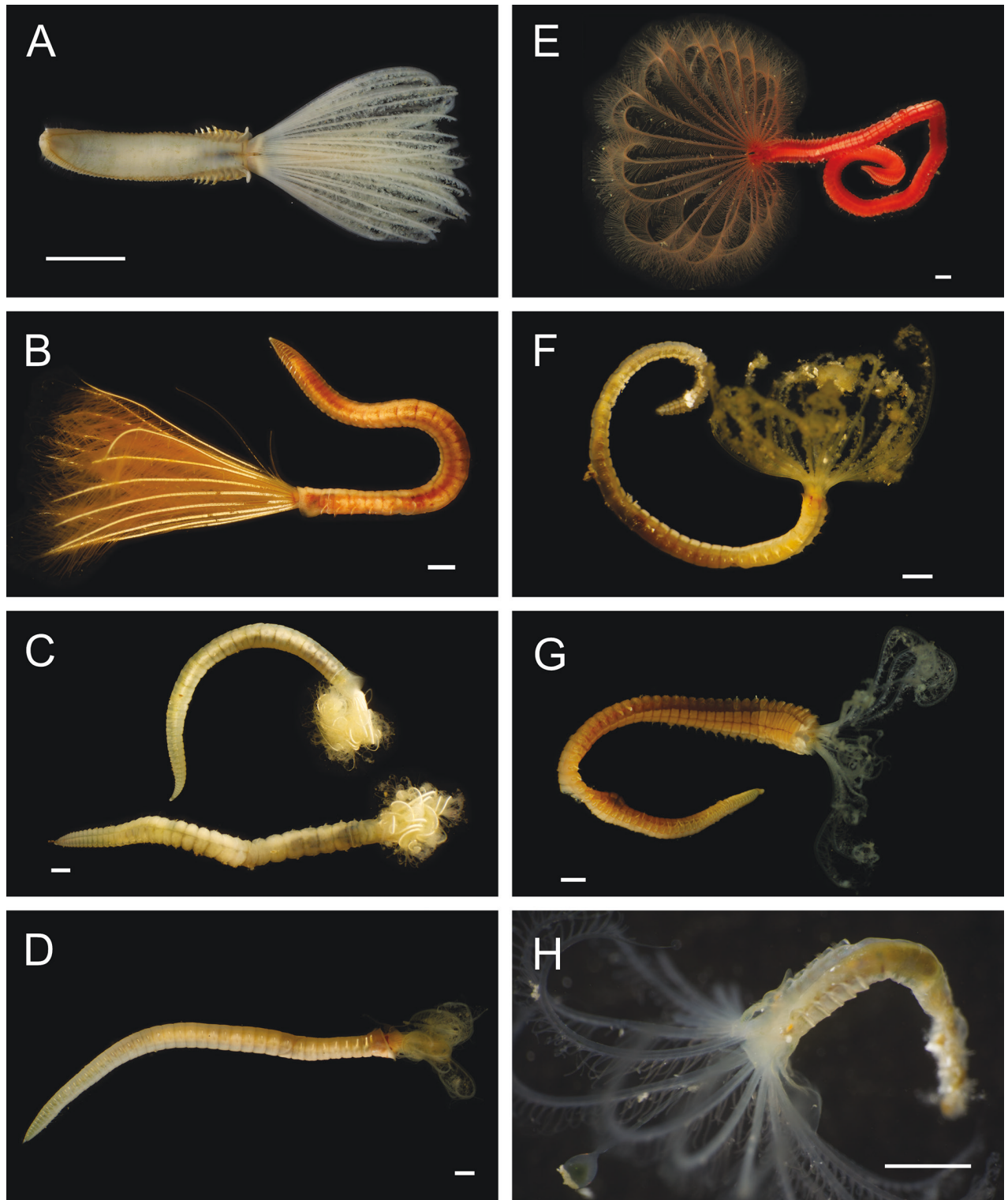


Figure 17. Annelida: Sabellidae and Serpulidae, representative live images **A** *Bispira* sp. SIO_BIC_A9587 (A9587) **B** *Chone* sp. SIO_BIC_A8422 (A8422) **C** *Chone* sp. SIO_BIC_A8462 (A8462) **D** *Jasmineira* stet. (A8282) **E** *Pseudopotamilla* sp. SIO_BIC_A1455 (A1455) **F** *Pseudopotamilla* sp. SIO_BIC_A9732 (A9732) **G** Sabellidae sp. SIO_BIC_ A8286 (A8286) **H** *Hyalopomatus* sp. SIO_BIC_A1434 (A1434). Scale bars: 1 cm (**A**); 1 mm (**B–H**).

Spirorbinae stet.

Fig. 18C

Material examined. AD4507: A1385.

Localities. Parrita Scar (~ 1659–1667 m).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Spionida |
Spionidae

***Aonides* sp. SIO_BIC_A1344**

Fig. 18D

Material examined. AD4503: A1344; AD4505: A1372, MZUCR 1509-01.

Localities. Mound 12 (~ 967–995 m), Mound 11 (~ 1019–1025 m).

Remarks. An undescribed species to be described in a forthcoming study.

***Lindaspio dibranchiata* Blake & Maciolek, 1992**

Fig. 18E

Material examined. AD4511: A1453 ([PQ449281](#)); AD4972: A9601 ([PQ449316](#));
S0213: A10053 ([PQ449255](#)).

Localities. Jacó Summit (~ 730–820 m), Mound 12 (989 m), Jacó Scar
(~ 1791–1800 m).

Distribution. Originally described from sedimented vents in the Guaymas
Basin, Gulf of California, 2004–2008 m (Blake and Maciolek 1992).

New records: The CRM specimens represent new southern records, the
first records from seeps, and, for specimen A10053, a new minimum depth for
this species (820 m as the most conservative value). The COI sequences for
the Costa Rica specimens were identical to that of a specimen from the type
locality: A3258 ([PQ432663](#)) from 1581 m at Pinkie's "Vent", Guaymas Basin.

***Prionospio* stet.**

Fig. 18F

Material examined. AD4509: A1415; AD4979: A9809.

Localities. Quepos Slide (393 m), Jacó Scar (1855 m).

Remarks. These specimens are members of the *Prionospio* complex;
the shape of the head of A1415 is consistent with *Apoprionospio*, but some
sources place this genus into synonymy with *Prionospio* (Vasily Rada-
shevsky, pers. comm. 8 July 2021).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Capitellida |
Opheliidae

Opheliidae stet.

Fig. 18G

Material examined. AD4971: A9651; AD4972: A9596; AD4977: A9730; AD4989:
A9938 ([PQ449337](#)).

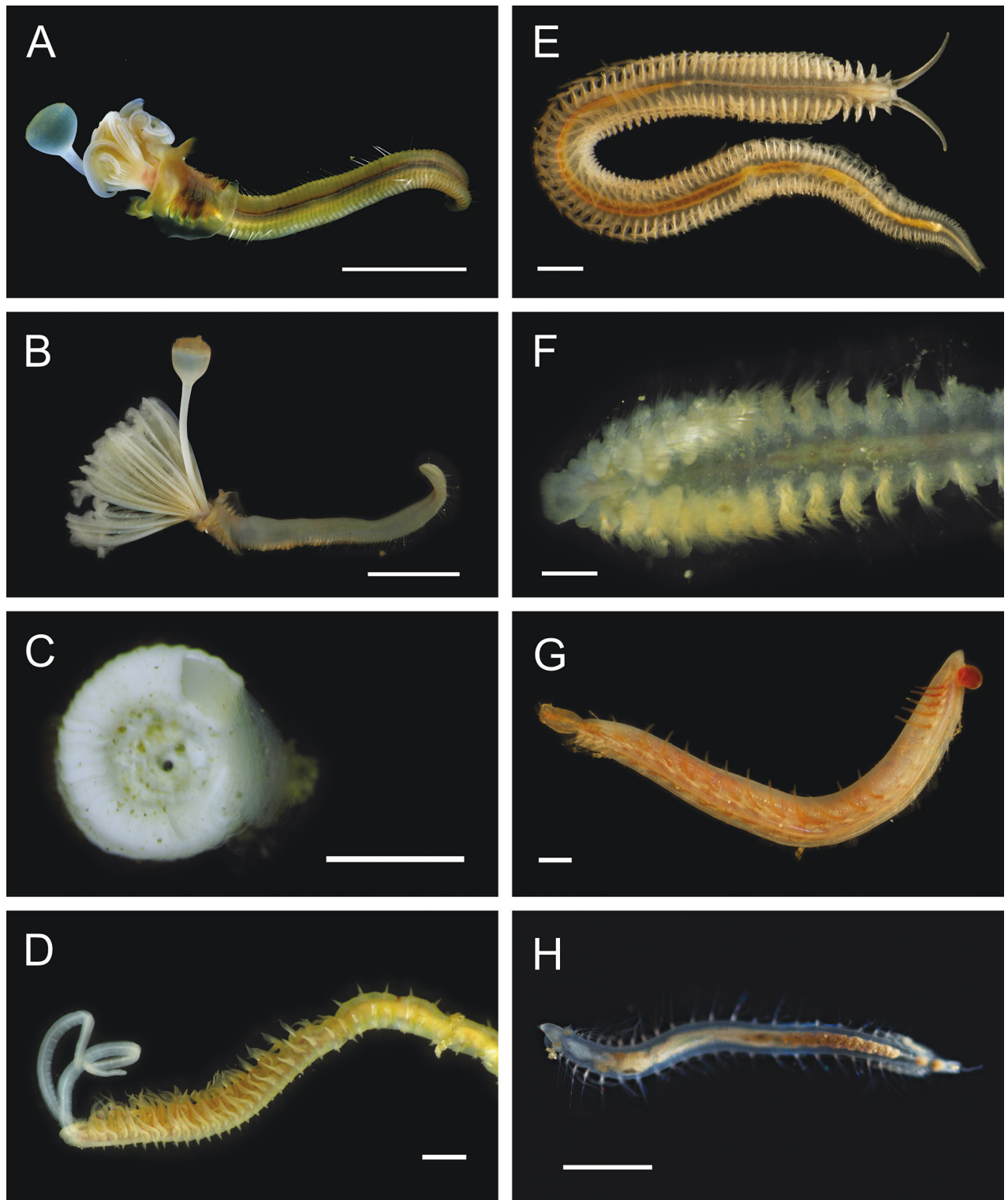


Figure 18. Annelida: Serpulidae, Spionidae, and Opheliidae, representative live images **A** *Laminatubus joycebrooksae* (A1315) **B** *Laminatubus paulbrooksi* (MZUCR 1506-01) **C** Spirobinae stet. (A1385, body not visible) **D** *Aonides* sp. SIO_BIC_A1344 (A1344) **E** *Lindaspio dibranchiata* (A10053) **F** *Prionospio* stet. (A1415) **G** Opheliidae stet. (A9938) **H** *Ophelina* sp. 3 sec. Law et al. 2014 (A1898). Scale bars: 1 cm (**A**, **B**); 1 mm (**C**–**H**).

Localities. Jacó Scar (1783–1796 m).

Remarks. A9651 likely represents an undescribed species of *Ophelina* (Sergio Salazar-Vallejo, pers. comm. 12 February 2020).

***Ophelina* sp. 3 sec. Law et al. 2014**

Fig. 18H

Reference. Law et al. (2014), in which GenBank sequences published under the temporary code F14588 ([KF511809](#), [KF511823](#), [KF511847](#), [KF511864](#)) correspond to specimen A1898 from AD4587.

Localities. Mound 12 (~ 990–996 m).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Capitellida | Capitelliformia | Thalassematidae

We refer to the clade formerly known as Echiura using the revised taxonomy of Goto et al. (2020).

***Prometor* stet.**

Fig. 19A

Material examined. AD4971: A9638 ([PQ449318](#)).

Localities. Jacó Scar (1824 m).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Capitellida | Capitelliformia | Capitellidae

***Capitellidae* stet.**

Fig. 19B

Material examined. AD4506: A1376; AD4508: A1403; AD4509: A1418; AD4976: A9755 (16S: [PQ304666](#)).

Localities. Parrita Seep (~ 1186–1419 m), Jacó Scar (~ 974–1887 m).

Remarks. Associated with vesicomyid clams (A1376, A1418), *Lamellibrachia barhami* (A1403), or experimentally deployed wood (A9755).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Terebellida | Scalibregmatidae

***Scalibregma* stet.**

Fig. 19C

Material examined. AD4507: A1391 ([PQ449279](#)); AD4914: A8385.

Localities. Parrita Scar (~ 1659–1667 m), Jacó Scar (1886 m).

***Travisia* stet.**

Fig. 19D

Material examined. AD4507: A1386.

Localities. Parrita Scar (~ 1659–1663 m).

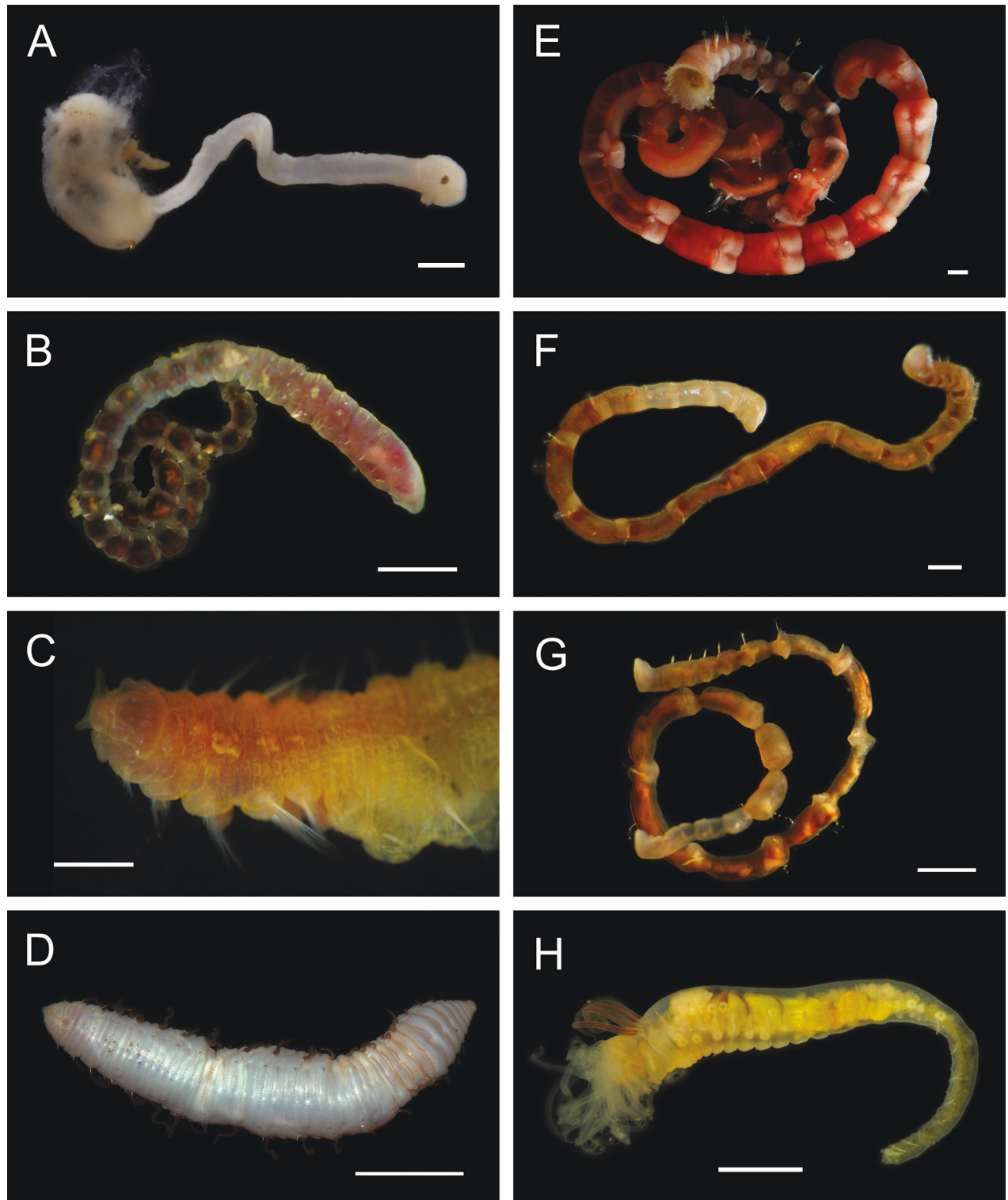


Figure 19. Annelida: Thalassematidae, Capitellidae, Scalibregmatidae, Maldanidae, and Ampharetidae, representative live images **A** *Prometor* stet. (A9638) **B** Capitellidae stet. (A1418) **C** *Scalibregma* stet. (A1391) **D** *Travisia* stet. (A1386) **E** *Nicomache* cf. *lokii* (A8257) **F** *Notoproctus* sp. SIO_BIC_A9801 (A9801) **G** *Notoproctus* sp. SIO_BIC_A9802 (A9802) **H** Ampharetini stet. (A1377). Scale bars: 1 mm (**A–C, E–H**); 1 cm (**D**).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Terebellida |
Maldanomorpha | Maldanidae

***Nicomache* cf. *lokii* Kongsrud & Rapp, 2012**

Fig. 19E

Material examined. AD4504: A1350 ([PQ450403](#)); AD4509: A1420 ([PQ450404](#));
AD4906: A8257; AD4911: A8295.

Localities. Mound 11 (~ 1002–1011 m), Mound 12 (995–1001 m), Jacó Scar
(~ 974–1856 m).

Remarks. The closest COI BLASTN results on GenBank were sequences
of *Nicomache lokii* (95.03–98.17% identity to A1350, 95.18–98.31% iden-
tity to A1420), and the closest matches ([MG975502.1](#), [FR877578.1](#)) were
from the type locality of the Loki's Castle vent system on the Arctic Mid-
Ocean Ridge at 2350 m (Kongsrud and Rapp 2012; Eilertsen et al. 2018).
Nicomache lokii shows genetic connectivity across vents and seeps in the
Arctic, Barbados Trench, and East Scotia Sea, 1262–4930 m, with a maxi-
mum intraspecific pairwise distance of 4.1% (K2P) (Eilertsen et al. 2018),
so further genetic and morphological comparisons to the CRM specimens
are warranted.

***Notoproctus* sp. SIO_BIC_A9801**

Fig. 19F

Material examined. AD4978: A9801 ([PQ449327](#)).

Localities. Mound 12 (997 m).

Remarks. This specimen showed a COI difference of 19.1% (uncorrected)
from A9802 (below) so we regard them as separate species.

***Notoproctus* sp. SIO_BIC_A9802**

Fig. 19G

Material examined. AD4978: A9802 ([PQ449328](#)).

Localities. Mound 12 (997 m).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Terebellida |
Terebelliformia | Ampharetidae

***Ampharetini* stet.**

Fig. 19H

Material examined. AD4506: A1377 ([PQ450382](#)).

Localities. Parrita Seep (1030 m).

***Amphisamytha fauchaldi* Solís-Weiss & Hernández-Alcántara, 1994**

Fig. 20A

References. Levin et al. 2012; Stiller et al. 2013.

Additional material examined. AD4906: A8260 ([PQ449288](#), [PQ449289](#)); AD4912: A8352 ([PQ449299](#), [PQ449300](#)); AD4913: A8358 ([PQ449302](#)); AD4914: A12602 ([PQ449273](#)).

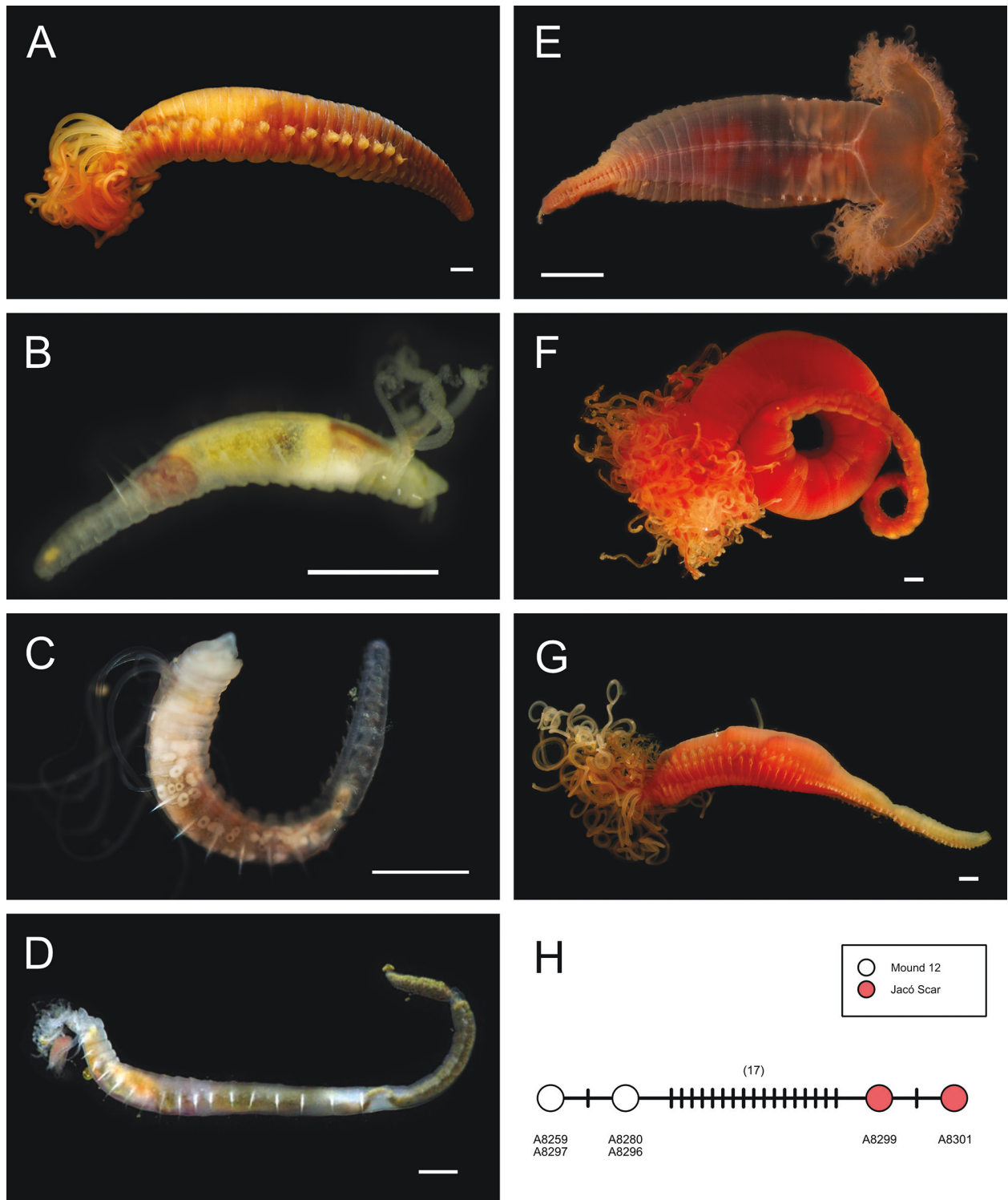


Figure 20. Annelida: Ampharetidae, Trichobranchidae, and Terebellidae, representative live images **A** *Amphisamytha fauchaldi* (A8352) **B** *Grassleia* cf. *hydrothermalis* (A12401) **C** *Pavelius* sp. EP-B sec. Stiller et al. 2020 (A1894) **D** *Terebellides* stet. (A1293) **E** *Biremis* sp. SIO_BIC_A10093 (A10093) **F** *Polycirrus* stet. (A10115) **G** *Eupolymnia* cf. *heterobranchia* (A8296) **H** Haplotype network of *Eupolymnia* cf. *heterobranchia* COI sequences. Scale bars: 1 mm (**A–D, F, G**); 1 cm (**E**).

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Distribution. Originally described from sedimented hydrothermal vents at 2000 m in the Guaymas Basin, Gulf of California (Solís-Weiss and Hernández-Alcántara 1994), *A. fauchaldi* is also found at seeps at Hydrate Ridge off Oregon, with an overall depth range of 600–2860 m (Stiller et al. 2013).

***Grassleia* cf. *hydrothermalis* Solís-Weiss, 1993**

Fig. 20B

Material examined. AD4510: A12401 ([PQ449272](#)).

Localities. Jacó Summit (744 m).

Remarks. The closest COI BLASTN result on GenBank was a specimen identified as *Grassleia* cf. *hydrothermalis* from seeps at 1572–1583 m in the Guaymas Basin, Gulf of California ([KX497032.1](#), 96.51% identity). *Grassleia hydrothermalis* was described from vents at the Escabana Trough, Gorda Ridge, off northern California, at 3271 m (Solís-Weiss 1993), and has been reported from seeps at 595 m off the Cascadia Margin (Reuscher et al. 2012). Genetic sampling from the type locality will be important to ascertain the distribution of this species.

***Pavelius* sp. EP-B sec. Stiller et al. 2020**

Fig. 20C

Material examined. AD4586: A1887 ([PQ449284](#)); AD4587: A1894 ([PQ449285](#)).

Localities. Mound 12 (~ 982–998 m).

Remarks. The CRM specimens represent records of an undescribed species also reported from seeps at Hydrate Ridge off Oregon, 809 m, as cited in supplementary table S2 of Stiller et al. (2020).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Terebellida |
Terebelliformia | Trichobranchidae

***Terebellides* stet.**

Fig. 20D

Material examined. AT15-44 MC1-2: A1293.

Localities. Near Mound 12 (1019 m).

Remarks. This specimen was collected in the upper 1 cm of a sediment core adjacent to Mound 12, ca 500 m from known sites of active seepage and likely representing the far-transition zone to the surrounding environment.

Annelida | Polychaeta | Sedentaria | Canalipalpata | Terebellida |
Terebelliformia | Terebellidae | Polycirri

***Biremis* sp. SIO_BIC_A10093**

Fig. 20E

Material examined. S0218: A10091, A10092, A10093.

Localities. Parrita Scar (1110–1470 m).

Remarks. An undescribed species.

Polycirrus stet.

Fig. 20F

Material examined. S0219: A10067, A10097, A10115 ([PQ449270](#)).

Localities. Rio Bongo Scar (609–659 m).

Remarks. These specimens were buried or partially buried in soft sediments with tentacles extended.

Annelida | Polychaeta | Sedentaria | Canalipalpata | Terebellida |
Terebelliformia | Terebellidae | Procleini

***Eupolymnia cf. heterobranchia* (Johnson, 1901)**

Fig. 20G

Material examined. AD4906: A8259 ([PQ449287](#)); AD4910: A8280 ([PQ449291](#)), A8296 ([PQ449294](#)), A8297 ([PQ450386](#)); AD4911: A8299 ([PQ449295](#)), A8301 ([PQ449296](#)).

Localities. Mound 12 (998–1002 m), Jacó Scar (1758 m).

Remarks. Likely an undescribed species, with morphological similarity to *Eupolymnia heterobranchia*, known from shallow waters from Alaska to Mexico (Banse 1980). The COI sequences showed low identity (~ 80–82%) to GenBank sequences of *E. heterobranchia* from the type locality of Puget Sound (Johnson 1901) ([HQ932678.1](#), [HM473379.1](#), [HM473380.1](#), [MN138388.1](#)). *Eupolymnia* is paraphyletic, and nomenclatural revision requires further phylogenetic sampling (Stiller et al. 2020).

The CRM specimens appeared to show genetic structure between Mound 12 and Jacó Scar, with 3.2–3.6% COI divergence (uncorrected, corresponding to 17–19 bp) between localities and a maximum divergence of only < 0.2% (1 bp) within localities (Fig. 20H). We conservatively consider the Mound 12 and Jacó Scar populations to represent the same taxon, pending further sampling and morphological investigation. Yet we acknowledge the possibility of depth-segregated cryptic species, given that the depth-spanning sympatric scaleworm sister species *Branchiopolynoe halliseyae* and *B. kajsae* are separated by a comparable COI distance (3.7% uncorrected) (Lindgren et al. 2019).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Terebellida |
Terebelliformia | Terebellidae | Terebellini

***Neoamphitrite cf. hydrothermalis* Reuscher, Fiege & Wehe, 2012**

Fig. 21A

Material examined. AD4504: A1351 ([PQ450387](#)).

Localities. Mound 11 (1010 m).

Remarks. Likely an undescribed species, morphologically similar to *Neoamphitrite hydrothermalis*, which is known from western Pacific hydrothermal vents in the Lihir Basin, 1474–1480 m (Reuscher et al. 2012). The COI sequence did not closely match any available GenBank reference sequences (<83% identity).

***Neoamphitrite* cf. *robusta* (Johnson, 1901)**

Fig. 21B

Material examined. AD4512: A1456 ([PQ450389](#)); AD4588: A2150 ([PQ449286](#)).

Localities. Quepos Slide (400 m), Mound 12 (~ 995–997 m).

Remarks. Likely an undescribed species. Morphologically similar to *Neoamphitrite robusta*, which was described from shallow water in Puget Sound (Johnson 1901) and occurs in the eastern Pacific to depths of at least 1984 m (Hartman and Barnard 1958), including Pacific Costa Rica at 22 m (Wehrtmann and Cortés 2009b). The COI sequences did not closely match any available GenBank reference sequences (<81% identity).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Terebellida |
Terebelliformia | Melinnidae

***Melinnopsis* cf. *armipotens* (Moore, 1923)**

Fig. 21C, D

Material examined. S0220: A12604 ([PQ449274](#)).

Localities. Subduction Plume (3502 m).

Remarks. The tubes of these animals protruded above the sediment and were encrusted with *Candelabrum* hydroids (Co3088). They warrant comparison to *Melinnopsis armipotens*, known only from southern California, 4075 m (Moore 1923).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Cirratulida |
Cirratulidae

We thank Jim Blake (Aquatic Research & Consulting) for morphological identification of these specimens. These specimens will be further discussed in a separate work.

***Aphelochaeta* sp. SIO_BIC_A1380**

Fig. 21E, F

Material examined. AD4506: A1380 ([PQ449000](#)), AD4510: A1429.

Localities. Jacó Summit (741 m), Parrita Seep (1186 m).

***Aphelochaeta* sp. SIO_BIC_A9729**

Fig. 21G

Material examined. AD4977: A9729 ([PQ449323](#)).

Localities. Jacó Scar (1783 m).

Remarks. Likely an undescribed species.

***Chaetozone* sp. SIO_BIC_A9846**

Fig. 21H

Material examined. AD4987: A9846 ([PQ449331](#)).

Localities. Mound 12 (999 m).

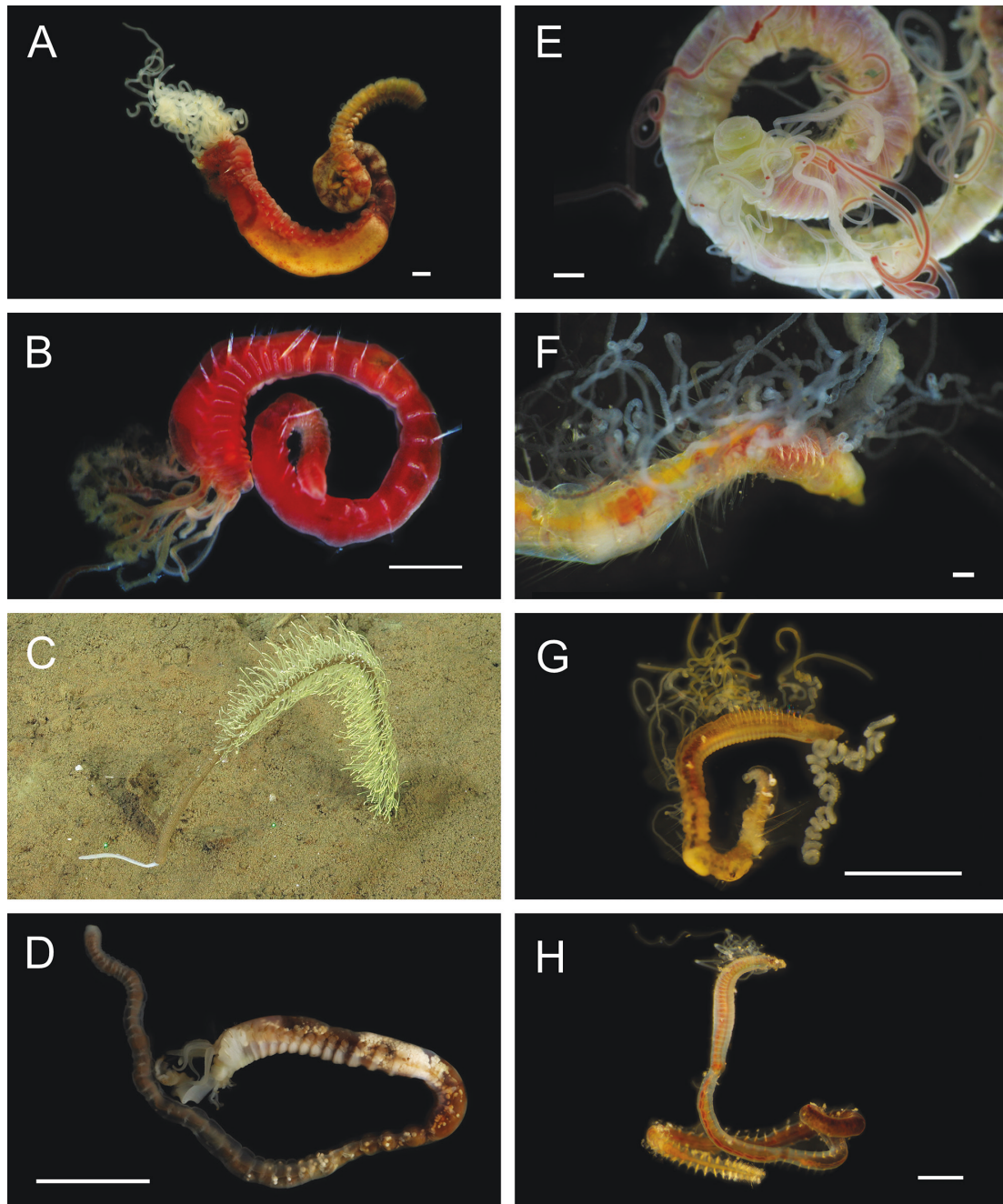


Figure 21. Annelida: Terebellidae, Melinnidae, and Cirratulidae, representative live images **A** *Neoamphitrite* cf. *hydrothermalis* (A1351) **B** *Neoamphitrite* cf. *robusta* (A1456) **C** *Melinnopsis* cf. *armipotens* (A12604, *in situ*, tentacle protruding from the tube extending from the sediment and covered with hydroids, Co3088). Credit: ROV SuBastian/Schmidt Ocean Institute **D** *Melinnopsis* cf. *armipotens* (A12604) **E** *Aphelochaeta* sp. SIO_BIC_A1380 (A1380) **F** *Aphelochaeta* sp. SIO_BIC_A1380 (A1429) **G** *Aphelochaeta* sp. SIO_BIC_A9729 (A9729) **H** *Chaetozone* sp. SIO_BIC_A9846 (A9846). Scale bars: 1 mm (**A**, **B**, **E–H**); 1 cm (**D**).

Remarks. An undescribed species.

***Cirratulus* stet.**

Fig. 22A

Material examined. AD4510: A1435 ([PQ449280](#)); AD4912: A8303 ([PQ449297](#)); AD4973: A9677 ([PQ449320](#)); AD4989: A9907 ([PQ449334](#)), A9962 ([PQ449339](#)); S0213: A10051 ([PQ449254](#)).

Localities. Jacó Summit (~ 741–744 m), Jacó Scar (~ 1768–1811 m).

Remarks. Multiple species may be represented.

***Raricirrus* cf. *maculatus* Hartman, 1961**

Fig. 22B

Material examined. AD4503: A1342 ([PQ449276](#)).

Localities. Mound 12 (990 m).

Remarks. Possibly *Raricirrus maculatus*, described from southern California, 46–70 m (Hartman 1961).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Cirratulida |
Acrocirridae

***Macrochaeta* stet.**

Fig. 22C

Material examined. AD4512: A1457; AD4918: A8420 ([PQ449307](#)); AD4985: A9854.

Localities. Quepos Slide (~ 338–411 m), Mound 12 (~ 995–1002 m).

Annelida | Polychaeta | Sedentaria | Canalipalpata | Cirratulida |
Flabelligeridae

We thank Sergio Salazar-Vallejo (El Colegio de la Frontera Sur) for morphological identification of these specimens.

***Bradabyssa* cf. *pilosa* (Moore, 1906)**

Fig. 22D

Material examined. AD4987: A9840 ([PQ449330](#)), A9902 ([PQ449333](#)).

Localities. Mound 12 (1010–1012 m).

Remarks. *B. pilosa* has been recorded from Alaska (type locality, 362–573 m) to Baja California, 40–1800 m, including seeps off Oregon and southern California (Salazar-Vallejo 2017). If confirmed as *B. pilosa* by genetic comparison to material from the type locality, the CRM specimens would represent a new southern record for the species. A9840 was associated with parasitic copepods (C14494), likely Bradophilidae.

***Bradabyssa* sp. SIO_BIC_A1356**

Fig. 22E

Material examined. AD4504: A1356 ([PQ449277](#)); AD4590: A1962 ([PQ450390](#)).

Localities. Mound 11 (~ 1004–1011 m), Jacó Scar (~ 1791–1800 m).

***Flabelligera* cf. *bophortica* Annenkova-Chlopina, 1924**

Fig. 22F

Material examined. AD4512: A1458; AD4976: A9751 ([PQ449324](#)).

Localities. Quepos Slide (~ 344–411 m), Jacó Scar (1887 m).

Remarks. These specimens resemble *F. bophortica*, which was described from the Chukchi Sea, Arctic Ocean, 13–48 m (Salazar-Vallejo 2012). The main morphological difference is that in *F. bophortica* the dorsum is areolate, whereas in the CRM specimens it is smooth (Sergio Salazar-Vallejo, pers. comm. 12 February 2020). Specimen A9751 was associated with experimentally deployed wood.

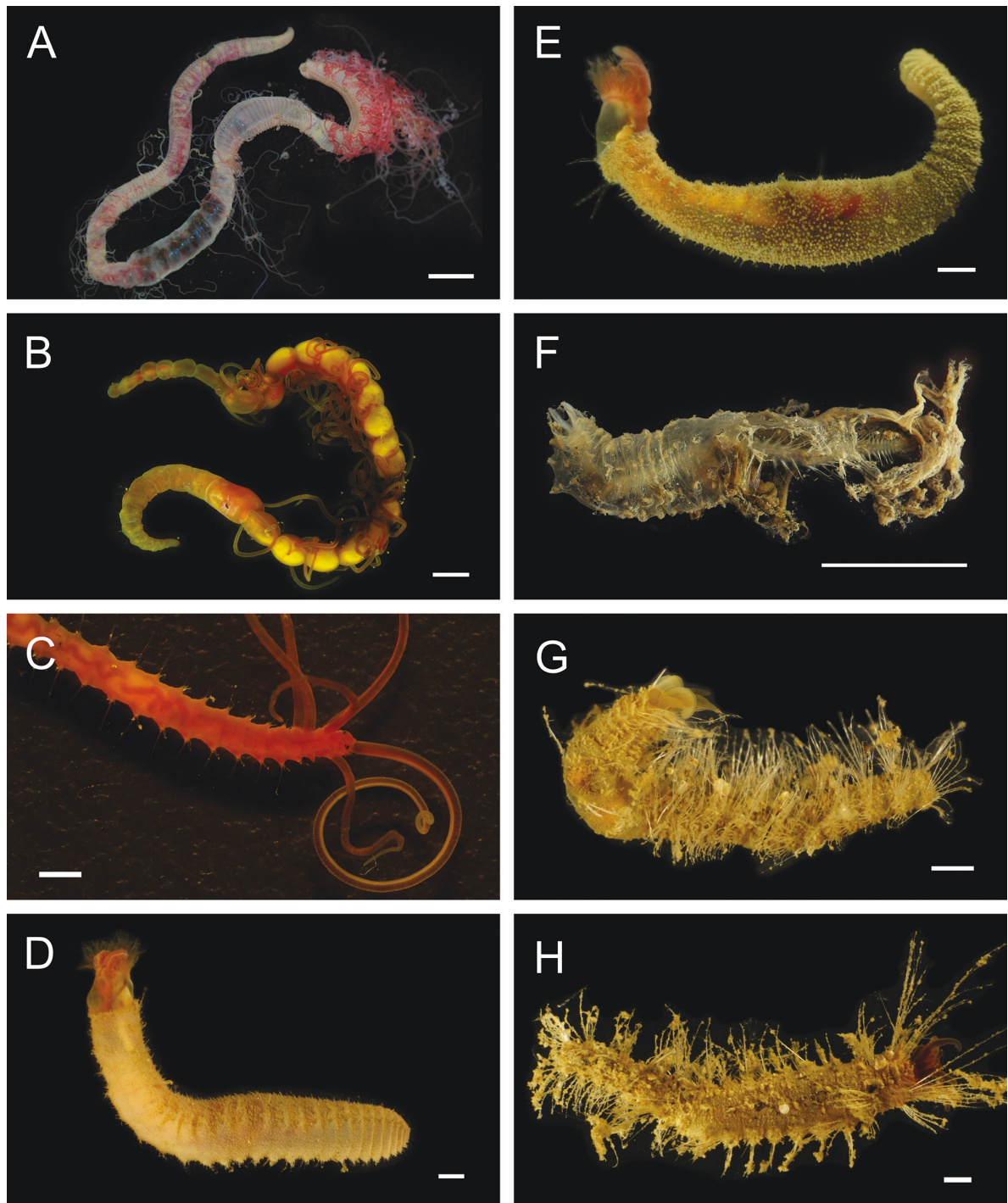


Figure 22. Annelida: Cirratulidae and Flabelligeridae, representative live images **A** *Cirratulus* stet. (A1435) **B** *Raricirrus* cf. *maculatus* (A1342) **C** *Macrochaeta* stet. (A8420) **D** *Bradabyssa* cf. *pilosa* (A9902) **E** *Bradabyssa* sp. SIO_BIC_A1356 (A1356) **F** *Flabelligera* cf. *bophortica* (A9751) **G** *Lamispina polycerata* (MZUCR 1504-01) **H** *Saphobranchia canela* (A9607). Scale bars: 1 cm (**A, F**); 1 mm (**B–E, G, H**).

***Lamispina polycerata* Salazar-Vallejo, 2020**

Fig. 22G

Reference. Salazar-Vallejo 2020a**.

Localities. Mound 12 (999 m; type locality).

Distribution. Known only from the CRM seeps.

***Saphobranchia canela* Salazar-Vallejo, 2020**

Fig. 22H

Reference. Salazar-Vallejo 2020a**.

Localities. Mound 12 (~ 987–997 m; type locality), Jacó Scar (1785 m).

Distribution. Known only from the CRM seeps.

Remarks. *S. canela* is distinguished from *S. ilys* and *S. omorpha* based on morphology, but genetic data do not corroborate the delineation of three species, as acknowledged in the original description (Salazar-Vallejo 2020a). The COI variation between specimens described as *S. ilys* and *S. omorpha* is less than the variation within *S. canela* (Fig. 23A). Collection of additional specimens and investigation of potential polymorphic conditions will be important for further work.

***Saphobranchia ilys* Salazar-Vallejo, 2020**

Fig. 23B

Reference. Salazar-Vallejo 2020a**.

Localities. Jacó Scar (1783–1784 m; type locality).

Distribution. Known only from the CRM seeps.

***Saphobranchia omorpha* Salazar-Vallejo, 2020**

Fig. 23C

Reference. Salazar-Vallejo 2020a**.

Localities. Jacó Scar (1795 m; type locality).

Distribution. Known only from the CRM seeps.

Annelida | Polychaeta | Sedentaria | Canalipalpata | Sternaspida | Paraonidae

***Aricidea mirifica* Strelzov, 1973**

Fig. 23D

Reference. Langeneck et al. (2019) for DNA sequences and phylogenetic analysis, in which GenBank sequences [MH700680](#) and [MH688926](#) correspond to specimen A1464.

Material examined. AT15-44 MC1-2: A1464.

Localities. Near Mound 12 (1019 m).

Distribution. Also reported from shallow Pacific waters of Costa Rica, 18–26 m, as well as the eastern and western Pacific and Antarctic (Wehrtmann and Cortés 2009b).

Remarks. This specimen was collected in a sediment core adjacent to Mound 12, ca 500 m from known sites of active seepage and likely representing the far-transition zone to the surrounding environment.

***Aricidea rubra* Hartman, 1963**

Fig. 23E

Reference. Langeneck et al. (2019) for DNA sequences and phylogenetic analysis, in which GenBank sequences [MH700679](#), [MH688925](#), and [MH700708](#) correspond to specimen A1616 from Mound 12.

Material examined. AD4511: A1616; S0213: A10049 ([PQ449252](#)).

Localities. Jacó Summit (~ 730–820 m; this study), Mound 12 (988 m).

Distribution. Originally described from submarine canyons in southern California, 603–1298 m (Hartman 1963) and recorded from the South Atlantic, eastern and western Pacific, and Scotia Sea (Blake 1996).

***Aricidea* sp. A sec. Langeneck et al. 2019**

Fig. 23F

Reference. Langeneck et al. (2019) for DNA sequences and phylogenetic analysis, in which GenBank sequences [MH700686](#), [MH688937](#), and [MH700724](#) correspond to specimen A1970.

Material examined. AD4591: A1970.

Localities. Jacó Scar (~ 1752–1795 m).

Remarks. This specimen is an epitoke.

Annelida | Polychaeta | Sedentaria | Canalipalpata | Sternaspida | Sternaspidae

***Sternaspis* stet.**

Fig. 23G

Reference. Drennan et al. (2019) for phylogenetic analysis.

Material examined. AT15-44 MC1-2: A1473 (16S: [MK810080](#); 18S: [MK809977](#)); AD4989: A9943, A9945 ([PQ449338](#)), A9948.

Localities. Near Mound 12 (1019 m), Jacó Scar (1762 m; this study).

Remarks. Based on 16S and 18S sequences, specimen A1473 is phylogenetically distinct from its closest reported relative, the Antarctic species *Sternaspis sendalli* Salazar-Vallejo, 2014 (Drennan et al. 2019). Consistent with this result, the closest COI BLASTN results on GenBank for A9945 were specimens of *S. sendalli* (e.g., [MK810006.1](#), [NC_068907.1](#); up to 90.85% identity). A1473 was collected in a sediment core adjacent to Mound 12, ca 500 m from known sites of active seepage and likely representing the far-transition zone to the surrounding environment.

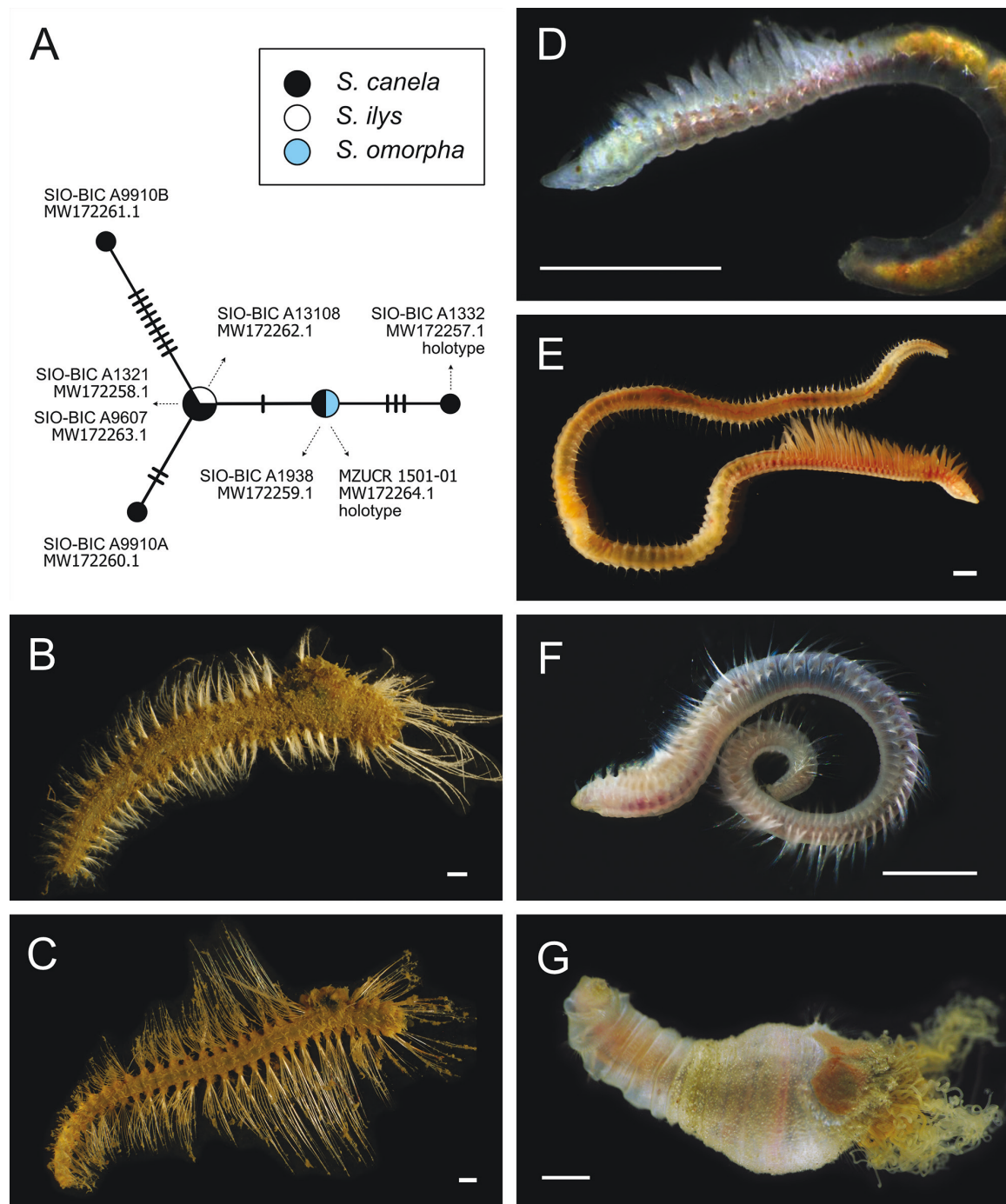


Figure 23. Annelida: Flabelligeridae, Paraonidae, and Sternaspidae, representative live images **A** haplotype network of COI sequences from the descriptions of *Saphobranchia canela*, *S. ilys*, and *S. omorpha* **B** *Saphobranchia ilys* (MZUCR 1503-01) **C** *Saphobranchia omorpha* (MZUCR 1501-01) **D** *Aricidea mirifica* (A1464) **E** *Aricidea rubra* (A10049) **F** *Aricidea* sp. A sec. Langeneck et al. 2019 (A1970) **G** *Sternaspis stet.* (A1473). Scale bars: 1 mm.

Annelida | Polychaeta | Sedentaria | Canalipalpata | Siboglinidae

***Escarpia spicata* Jones, 1985**

Fig. 24A

References. Levin et al. (2012, 2015) for occurrences at Mound 12, Jacó Scar, and Parrita Seep.

Material examined. AD4503: A1343; AD4507: A1390 ([PQ450388](#)); AD4511: A1445; AD4513: A1463; AD4590: A1826; AD4591: A1838; AD4914: A8364; AD4924: A8464; AD4971: A9618, A9619, A9620, A9621, A9622, A9623, A9624, A9625, A9626, A9627; AD4987: A9901; S0217: A10089; S0230: A10172.

Localities. Mound 12 (~ 1000 m), The Thumb (1074 m; this study), Parrita Seep (~ 1400 m), Parrita Scar (~ 1600 m; this study), Jacó Scar (~ 1800 m), Mound Jaguar (1909 m; this study).

Distribution. Originally described from seeps at 1829 m in the San Clemente Basin off California (Karaseva et al. 2016), *E. spicata* has been reported as far south as northern Chile (Kobayashi and Araya 2018). It has been recorded from a range of reducing environments and depths, including a whale fall at 1240 m near the type locality (Feldman et al. 1998); sedimented vents and seeps at 1568–2020 m in the Guaymas Basin, Gulf of California (Black et al. 1997; Karaseva et al. 2016); and a seep at 2756 m in the Middle America Trench, off southern Mexico (Southward et al. 2002).

Remarks. The three described species of *Escarpia* (*E. laminata* Jones, 1985; *E. southwardae* Andersen et al., 2004; *E. spicata* Jones, 1985) are morphologically distinguishable with separate geographic ranges (Karaseva et al. 2016), but the most commonly used DNA sequence markers (COI, 16S, CytB) do not provide reliable differentiation among these species due to low sequence variation (Black et al. 1997; McMullin et al. 2003; Cowart et al. 2013). Alternative marker sequences, such as the nuclear hemoglobin subunit B2 intron (HbB2i), have been used to support the existing species designations, but resolving their phylogenetic relationships remains complex (Cowart et al. 2013; Kobayashi and Araya 2018). Nevertheless, all eastern Pacific occurrences of *Escarpia* are presently accepted as *E. spicata* (Karaseva et al. 2016).

***Lamellibrachia barhami* Webb, 1969**

Fig. 24B

References. McMullin et al. 2003; Han et al. 2004; Mau et al. 2006; Sahling et al. 2008; Levin et al. 2012, 2015; McCowin and Rouse 2018.

Localities. Parrita Seep (~ 1400 m), Jacó Scar (~ 1800–1890 m), Parrita Scar (~ 2200 m).

Distribution. Originally described from seeps at 1125 m off southern California (Webb 1969), *L. barhami* occurs at seeps and sedimented vents (Middle Valley, Juan de Fuca Ridge; Pinkie's "Vent," Guaymas Basin), from Vancouver Island, Canada, to northern Chile, 1000–2400 m (Black et al. 1997, 1998; Bright and Lallier 2010; Karaseva et al. 2016; Kobayashi and Araya 2018; McCowin and Rouse 2018). The species shows only minimal COI divergence (<0.2%) across its range of more than 10,000 km (Kobayashi and Araya 2018; McCowin and Rouse 2018).

***Lamellibrachia donwalshi* McCowin & Rouse, 2018**

Fig. 24C

Reference. McCowin and Rouse 2018**.

Localities. Mound 12 (~ 1000 m; type locality), Mound 11 (~ 1040 m).

Distribution. Known only from the CRM seeps.

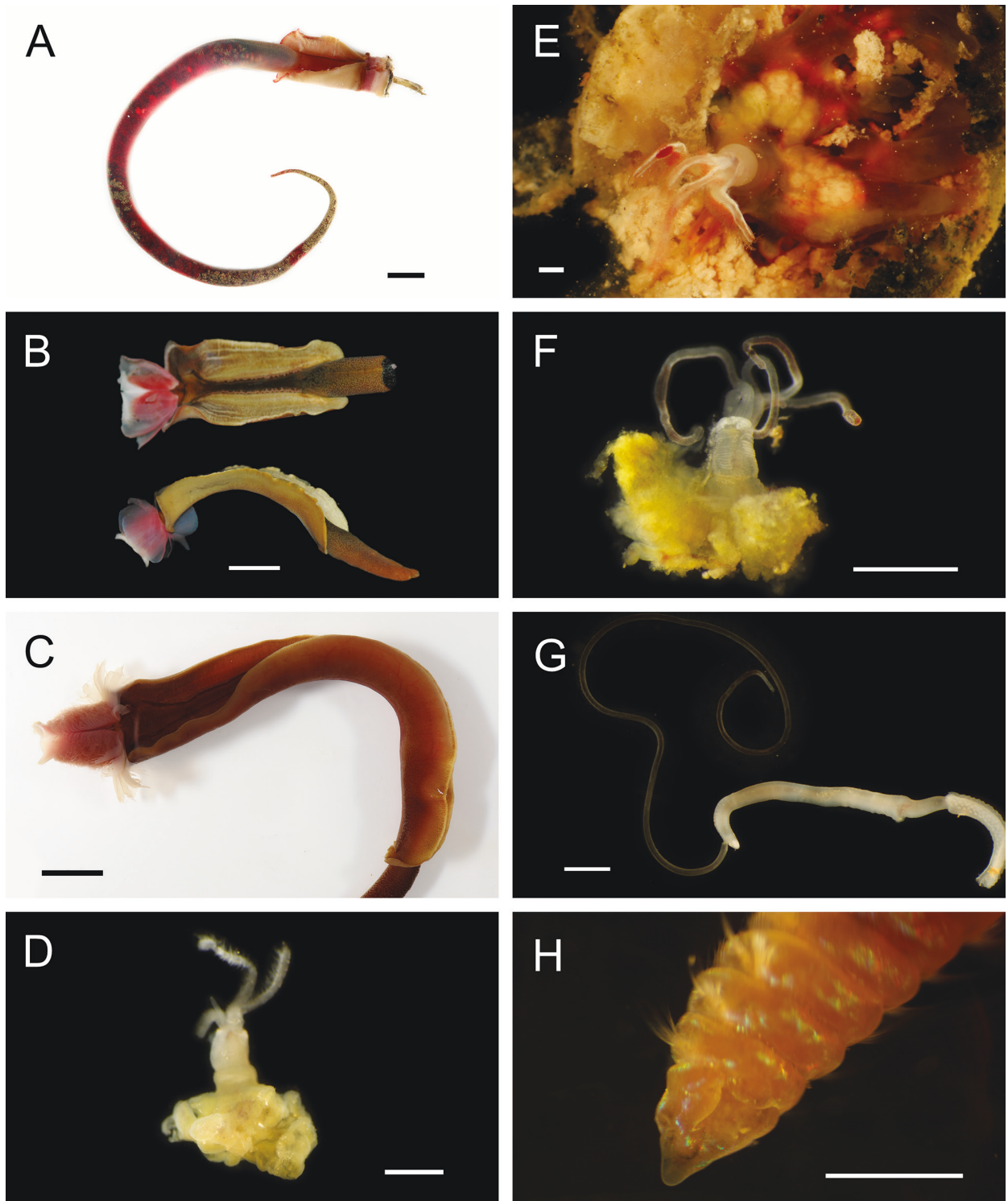


Figure 24. Annelida: Siboglinidae and Orbiniidae, representative live images **A** *Escarpia spicata* (A8364) **B** *Lamellibrachia barhami* (A1564, incomplete specimens) **C** *Lamellibrachia donwalshi* (A8382) **D** *Osedax frankpressi* (A9592) **E** *Osedax frankpressi* (A9594, in bone) **F** *Osedax knutei* (A9617) **G** *Siboglinum stet.* (A8349) **H** *Leitoscoloplos* sp. SIO_BIC_A9939 (A9939). Scale bars: 1 cm (**A–C**); 1 mm (**D–H**).

***Osedax frankpressi* Rouse, Goffredi & Vrijenhoek, 2004**

Fig. 24D, E

Reference. Berman et al. (2023) for DNA sequences and haplotype networks.

Material examined. AD4972: A9591 ([OM994442](#)), A9592 ([OM994444](#)), A9593 ([OM994443](#)), A9594 ([OM994445](#)).

Localities. Jacó Scar (1845 m).

Distribution. Originally described from Monterey Submarine Canyon off California (Rouse et al. 2004). Recorded in the eastern Pacific from the Oregon margin to the CRM as well as in the Atlantic off Brazil, 642–2891 m (Berman et al. 2023).

Remarks. Collected from experimentally deployed pig bones.

***Osedax knutei* Rouse, Goffredi, Johnson & Vrijenhoek, 2018**

Fig. 24F

Reference. Berman et al. (2023) for DNA sequences and haplotype networks.

Material examined. AD4974: A9617 ([ON041090](#)).

Localities. Mound 12 (992 m).

Distribution. Monterey Submarine Canyon off California (type locality) to the CRM, 845–2898 m (Berman et al. 2023).

Remarks. Collected from experimentally deployed pig or cow bones.

Siboglinum stet.

Fig. 24G

Material examined. AD4911: A8349 ([PQ449298](#)); AD4916: A8388; AD4989: A9942.

Localities. Jacó Scar (~ 1757–1892 m).

Remarks. Collected from sediment cores, except for A8349 which was associated with a rock substrate. The occurrence of a field of frenulate siboglinids at Jacó Scar was noted by Levin et al. (2012).

Annelida | Polychaeta | Sedentaria | Orbiniida | Orbiniidae

***Leitoscoloplos* sp. SIO_BIC_A9939**

Fig. 24H

Material examined. AD4989: A9939.

Localities. Jacó Scar (1785 m).

Remarks. This single damaged specimen likely represents an undescribed species (Jim Blake, pers. comm. 7 October 2019).

Annelida | Polychaeta | Sedentaria incertae sedis

***Cossura* stet.**

Fig. 25A

Material examined. AD4507: A1388; AD4988: A9922.

Localities. Mound 11 (~ 1005–1025 m), Parrita Scar (~ 1659–1663 m).

Annelida | Polychaeta | Amphinomida

Amphinomidae stet.

Fig. 25B

Material examined. S0219: A10107 ([PQ449269](#)).

Localities. Rio Bongo Scar (661 m).

Remarks. The COI sequence did not closely match any available GenBank reference sequences (<82% identity).

Amphinominae sp. SIO_BIC_A1379

Fig. 25C

Material examined. AD4506: A1379 ([PQ300673](#)).

Localities. Parrita Seep (1186 m).

Remarks. An undescribed genus and species. We thank Liz Borda (Texas A&M University San Antonio) for providing the COI sequence.

***Archinome levinae* Borda, Kudenov, Chevaldonné, Blake, Desbruyères, Fabri, Hourdez, Pleijel, Shank, Wilson, Schulze & Rouse, 2013**

Fig. 25D

Reference. Borda et al. 2013**.

Localities. Mound 12 (~ 1000 m), Mound 11 (~ 1040 m; type locality), Parrita Seep (1402 m), Jacó Scar (~ 1800 m).

Distribution. Also known from vents in the Guaymas Basin, Gulf of California, ~ 2400 m (Borda et al. 2013).

Annelida | Polychaeta | Sipuncula

Sipuncula sp. SIO_BIC_A9803

Fig. 25E

Material examined. AD4978: A9803 ([PQ449329](#)).

Localities. Mound 12 (997 m).

Remarks. The closest COI BLASTN results on GenBank were an unidentified sipunculan from southern California ([MK550656.1](#), 85.74% identity) and several golfingiids (~ 79–81% identity).

Sipuncula sp. SIO_BIC_A9839

Fig. 25F

Material examined. AD4987: A9839 ([PQ450416](#)).

Localities. Mound 12 (1012 m).

Remarks. Collected near a naturally occurring wood fall. The closest COI BLASTN results on GenBank were *Nephasoma abyssorum* (Koren & Danielssen, 1876) ([JN865109.1](#), 82.14% identity) and several other golfingiids (~ 78–79% identity).

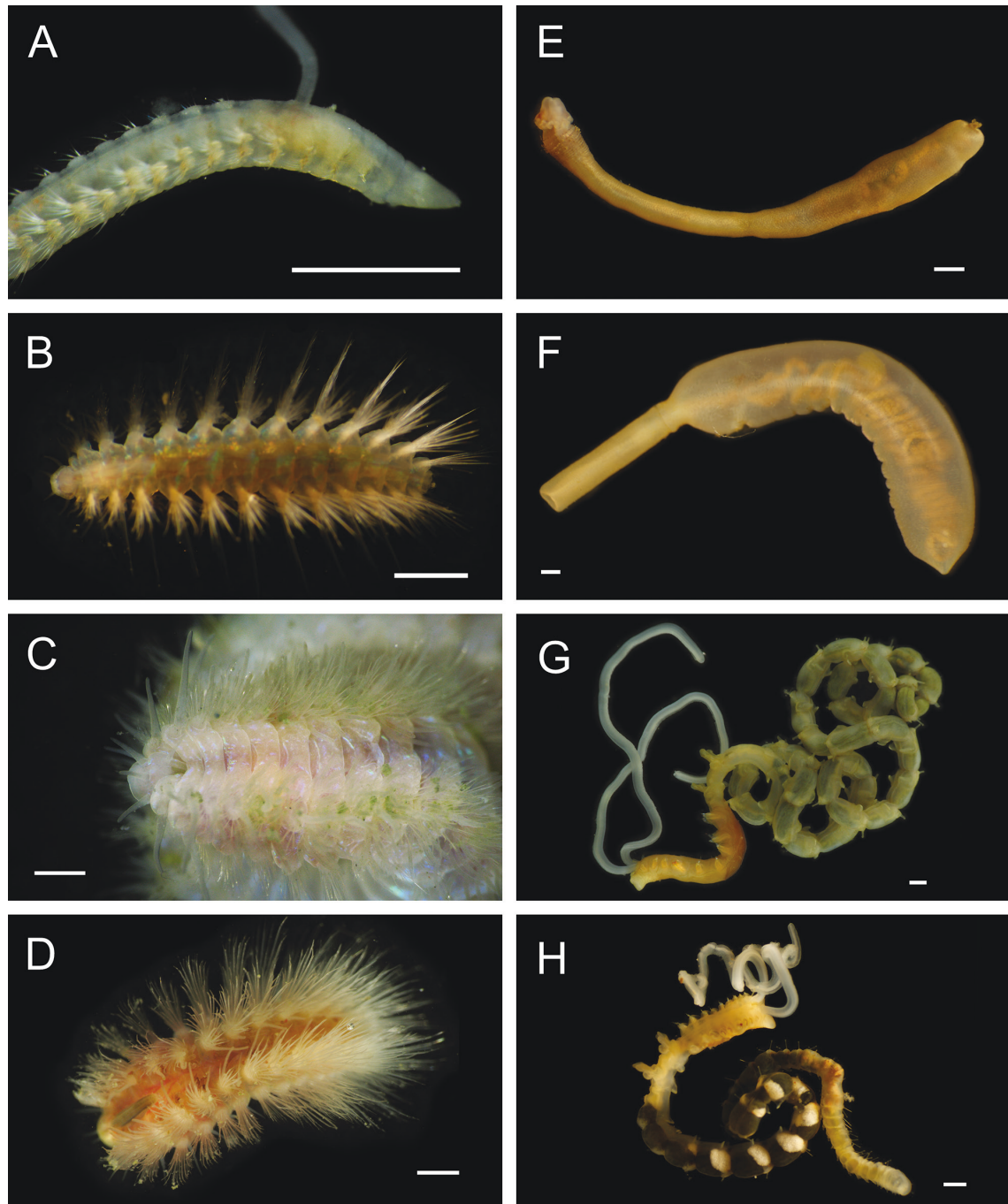


Figure 25. Annelida: *Cossura*, Amphinomidae, Sipuncula, and Chaetopteridae, representative live images **A** *Cossura* stet. (A1388) **B** Amphinomidae stet. (A10107) **C** Amphinominae sp. SIO_BIC_A1379 (A1379) **D** *Archinome leviniae* (A1398) **E** Sipuncula sp. SIO_BIC_A9803 (A9803) **F** Sipuncula sp. SIO_BIC_A9839 (A9839) **G** *Phyllochaetopterus* sp. 6 sec. Moore et al. 2017 (A1540) **H** *Phyllochaetopterus* sp. SIO_BIC_A8429 (A8429). Scale bars: 1 mm.

Annelida | Chaetopteriformia | Chaetopteridae

Phyllochaetopterus sp. 6 sec. Moore et al. 2017

Fig. 25G

Reference. Moore et al. (2017) for DNA sequences and phylogenetic analysis of specimen A1540.

Material examined. AD4501: A1324 ([PQ449275](#)); AD4505: A1540 ([KX896487](#)); AD4508: A1394; S0217: A10085 ([PQ449263](#)).

Localities. Mound 11 (~ 1019–1025 m), Mound 12 (~ 984–997 m; this study), The Thumb (~ 940–1070 m; this study), Parrita Seep (1402 m; this study).

Remarks. An undescribed species.

***Phyllochaetopterus* sp. SIO_BIC_A8429**

Fig. 25H

Material examined. AD4922: A8429 ([PQ449310](#)); AD4988: A9916 ([PQ449335](#)), A9918 ([PQ449336](#)).

Localities. Mound 11 (1010 m), Mound 12 (1002 m).

Remarks. Associated with naturally occurring wood falls. Despite its sympatry with *Phyllochaetopterus* sp. 6 at Mound 12, this undescribed species is morphologically and genetically distinct, with a COI divergence of 14.89–15.41% from the previously published *P.* sp. 6 voucher SIO-BIC A1540 ([KX896487.1](#)). The closest COI BLASTN result on GenBank was an undescribed species of *Spiochaetopterus* from the East Pacific Rise ([KX896501.1](#), voucher SIO-BIC A3619, 90.36–90.73% identity) (Moore et al. 2017). *Phyllochaetopterus* and *Spiochaetopterus* are not reciprocally monophyletic, and further phylogenetic investigation of their respective type species is required for revision of these genera (Moore et al. 2017).

Nemertea

We list higher-level taxonomy according to Strand et al. (2018).

Nemertea | Palaeonemertea | Tubulanidae

***Tubulanus* cf. *lutescens* Cantell, 2001**

Fig. 26A

Reference. Sagorny et al. 2022.

Localities. Mound 11 (~ 1019–1025 m), Mound 12 (~ 982–998 m).

Remarks. Despite disparate geography and internal and external morphological differences, the CRM specimens show only 1.3% COI divergence from *Tubulanus lutescens* Cantell, 2001, known from shallow waters off Sweden, warranting further examination (Sagorny et al. 2022).

Nemertea | Pilidiophora | Heteronemertea | Lineidae

***Lineidae* sp. SIO_BIC_N254**

Fig. 26B

Reference. Sagorny et al. 2022.

Localities. Jacó Scar (1887 m).

Remarks. Associated with experimentally deployed wood. This single specimen (destroyed for DNA extraction) could not be attributed to a known genus and likely represents an undescribed species (Sagorny et al. 2022).

Nemertea | Hoplonemertea | Monostilifera | Amphiporina

***Tetrastemma polyakovae* Sagorny, von Döhren, Rouse & Tilic, 2022**

Fig. 26C

Reference. Sagorny et al. 2022**.

Localities. Mound 12 (~ 996–999 m; type locality).

Distribution. Known only from the CRM seeps.

***Tetrastemma strandae* Sagorny, von Döhren, Rouse & Tilic, 2022**

Reference. Sagorny et al. 2022** (histological sections published; no live images captured).

Localities. Jacó Scar (1885 m; type locality).

Distribution. Known only from the CRM seeps.

***Tetrastemma sundbergi* Sagorny, von Döhren, Rouse & Tilic, 2022**

Fig. 26D

Reference. Sagorny et al. 2022**.

Localities. Mound 12 (~ 996–999 m; type locality).

Distribution. Known only from the CRM seeps.

Nemertea | Hoplonemertea | Monostilifera | Eumonostilifera

***Eumonostilifera* sp. SIO_BIC_N109**

Fig. 26E

Reference. Sagorny et al. 2022.

Localities. Parrita Seep (~ 1401–1419 m).

Remarks. An undescribed species represented by a single specimen (Sagorny et al. 2022).

Nemertea | Hoplonemertea | Monostilifera | Oerstediiina

***Alvinonemertes christianeae* Sagorny, von Döhren, Rouse & Tilic, 2022**

Fig. 26F

Reference. Sagorny et al. 2022**.

Localities. Jacó Scar (1887 m; type locality).

Distribution. Also reported from the non-seep seamount Quepos Plateau, 67 km south of Jacó Scar, at 2184 m depth.

Remarks. Associated with experimentally deployed or naturally occurring wood (Sagorny et al. 2022).

***Alvinonemertes dariae* Sagorny, von Döhren, Rouse & Tilic, 2022**

Fig. 26G

Reference. Sagorny et al. 2022**.

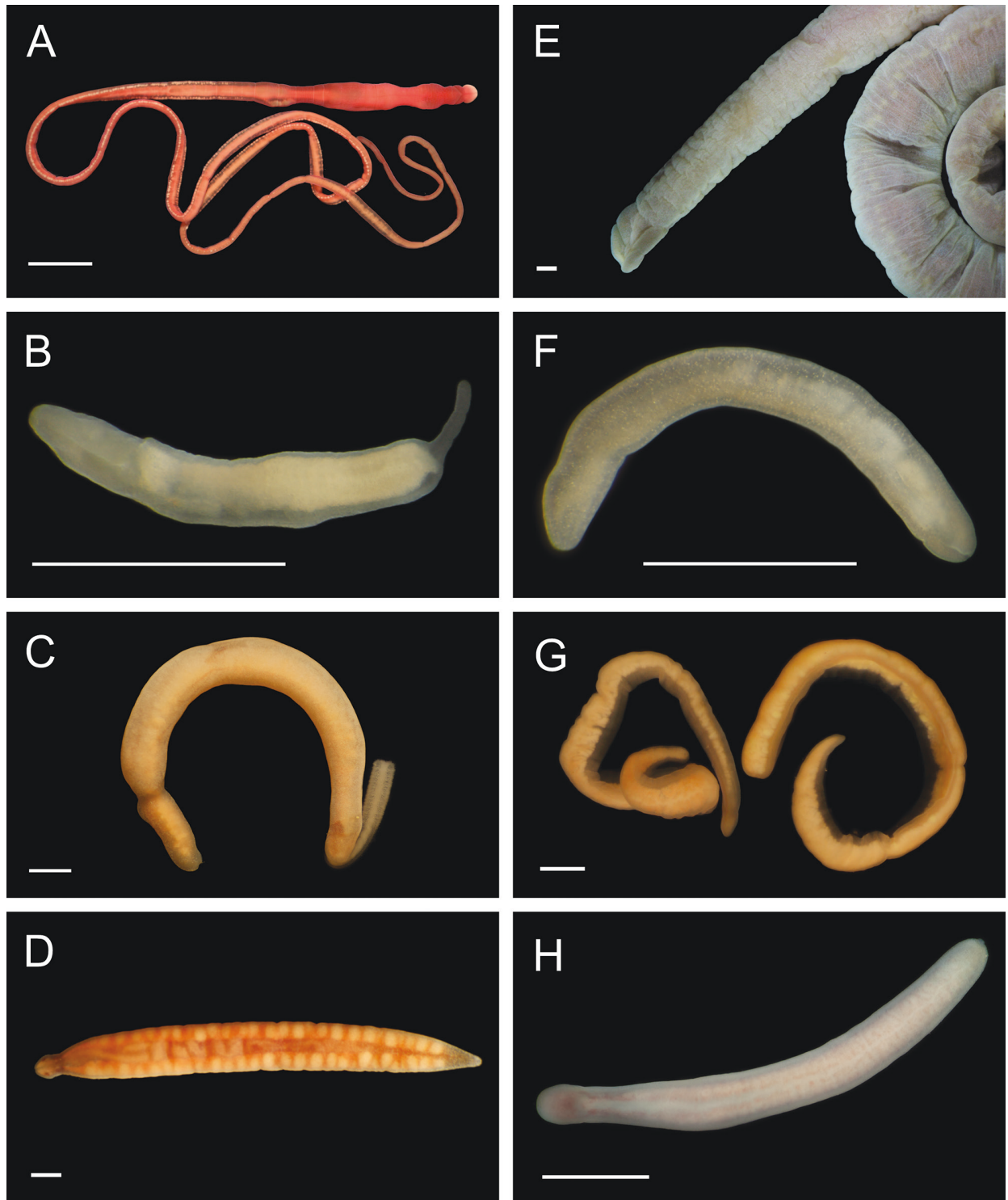


Figure 26. Nemertea, representative live images **A** *Tubulanus* cf. *lutescens* (N233) **B** Lineidae sp. SIO_BIC_N254 (N254) **C** *Tetrastemma polyakovae* (N258) **D** *Tetrastemma sundbergi* (N256) **E** *Eumonostilifera* sp. SIO_BIC_N109 (N109) **F** *Alvinonemertes christianeae* (N253) **G** *Alvinonemertes dariae* (N262) **H** *Chernyshevia escarpiaphila* (N266). Scale bars: 1 cm (A); 1 mm (B–H).

Localities. Parrita Seep (1407 m; type locality).

Distribution. Known only from the CRM seeps.

Remarks. Associated with *Paragorgia* stet. (Co3054) (Sagorny et al. 2022).

***Chernyshevia escarpiaphila* Sagorny, von Döhren, Rouse & Tilic, 2022**

Fig. 26H

Reference. Sagorny et al. 2022**.

Localities. Jacó Scar (~ 974–1856 m), Mound Jaguar (1909 m; type locality).

Distribution. Known only from the CRM seeps.

Remarks. Associated with the exterior surfaces and tubes of *Escarpia spicata* (Sagorny et al. 2022).

Brachiopoda

***Platidia anomioides* (Scacchi & Philippi in Philippi, 1844) sp. inc.**

Fig. 27A, B

Material examined. AD4512: B211; AD4979: M16876.

Localities. Quepos Slide (~ 380–395 m).

Remarks. We thank Sandra Carlson (University of California Davis) for the identification of these specimens as *Platidia*, most likely *P. anomioides*, which is considered cosmopolitan at depths 18–2190 m (Foster 1989; Santagata and Tunnell 2009). More definitive identification requires dissolving soft tissues for analysis of the cardinalia and spicules. At least one specimen contains eggs in the mantle canals (Fig. 27B).

Mollusca

We list the major clades according to the phylogenetic relationships in Smith et al. (2011).

Mollusca | Bivalvia

We list entries following the taxonomic arrangement in Valentich-Scott et al. (2020).

Mollusca | Bivalvia | Protobranchia | Nuculida | Nuculidae

***Ennucula colombiana* (Dall, 1908)**

Fig. 27C

Material examined. AD4588: M17879.

Localities. Mound 12 (997 m).

Distribution. Originally described from the Gulf of Panama, 54 m, and known from the Gulf of California to Peru, 11–734 m (Coan and Valentich-Scott 2012).

New records. The CRM specimen represents a new maximum depth record for this species.

***Ennucula* stet.**

Fig. 27D

Material examined. AD4988: M16966 ([PQ449418](#)).

Localities. Mound 11 (1010 m).

Remarks. This damaged juvenile specimen was associated with a naturally occurring wood fall. The closest COI BLASTN results on GenBank were nucleotides: *Ennucula cumingii* (Hinds, 1843) ([KC984750.1](#), 84.89% identity), *E. tenuis* (Montagu, 1808) from Japan ([LC144804.1](#), 83.23% identity), *E. niponica* (E. A. Smith, 1885) from Japan ([LC144803.1](#), 83.10% identity), and *Acila mirabilis* (A. Adams & Reeve, 1850) from Japan ([LC144802.1](#), 83.36% identity).

***Nucula chrysocoma* Dall, 1908**

Fig. 27E

Material examined. AD4503: M11989.

Localities. Mound 12 (~ 965–995 m).

Distribution. Originally described from several stations off Peru, Ecuador, and southern Mexico, 734–4064 m (Dall, 1908), and known from Cascadia Abyssal Plain, Oregon, to Coquimbo, central Chile, 734–4134 m (Valentich-Scott et al. 2020).

Mollusca | Bivalvia | Protobranchia | Solemyida | Solemyidae

***Acharax cf. johnsoni* (Dall, 1891)**

Fig. 27F, G

References. Neulinger et al. (2006) for DNA sequences and phylogenetic analysis; occurrences at Mound 11 and Mound 12 (Aguado and Rouse 2011; Levin et al. 2015; Rouse et al. 2018).

Additional material examined. AD4503: M11980; AD4505: M12003; AD4507: M12012 (juvenile); AD4511: M12054, M16239; AD4513: M12072; AD4910: M15768 (18S: [PQ304648](#)), M15769; S0217: M17062; S0220: M17077 (18S: [PQ304649](#); juvenile).

Localities. Mound 12 (~ 1000 m), Mound 11 (~ 1020 m), The Thumb (1069 m; this study), Parrita Scar (~ 1659–1663 m; this study), Jacó Scar (1744 m; this study), Subduction Plume (3410 m; this study). The collection locality in Neulinger et al. (2006) (9°10.4'N, 084°48.2'W, 763 m) matches the site referenced in this work as Jacó Summit.

Remarks. Originally described from 1838 m off Baja California (Dall 1891), *A. johnsoni* has been reported along the eastern Pacific margin from Alaska to Peru, as well as in the northwestern Pacific, 100–5379 m (Coan and Valentich-Scott 2012). According to 18S phylogenies, at least two clades of deep-sea *Acharax*, morphologically similar to *A. johnsoni*, occur in the Pacific and warrant consideration as separate species (Neulinger et al. 2006; Fukasawa et al. 2017). In the absence of genetic data from the type locality (only empty shells have been recovered from recent expeditions to the region (Hendrickx et al. 2016; Suárez-Mozo et al. 2019)), we report the CRM specimens as *A. cf. johnsoni*.

The two 18S sequences in this study matched different previously reported clades, suggesting the presence of at least two cryptic species at the CRM, potentially segregated by depth. M17007 from Subduction Plume (3410 m) showed 99.94% identity to the Jacó Summit specimen ([AJ563763.1](#)) in Neulinger et al.

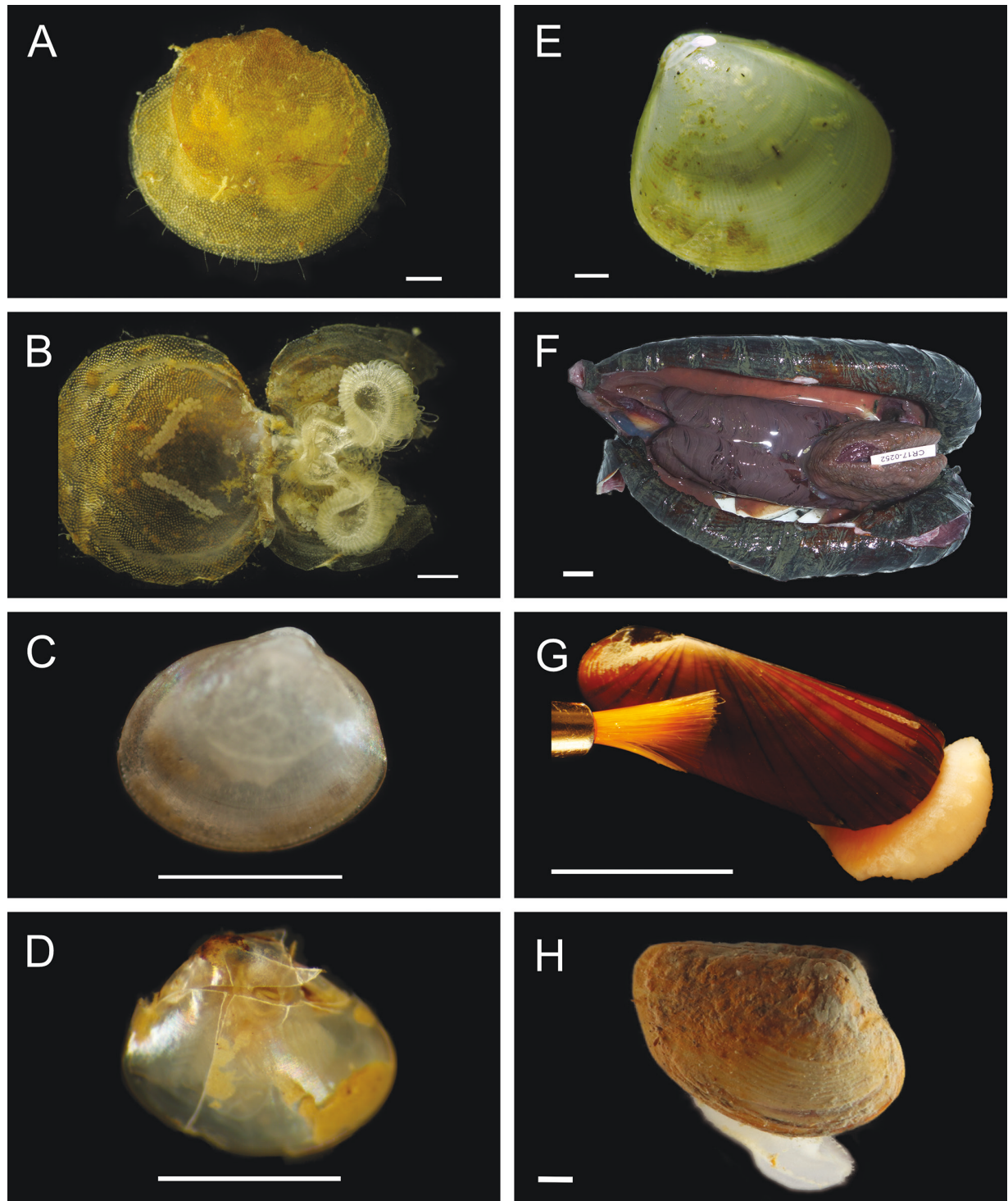


Figure 27. Brachiopoda and Mollusca: Bivalvia: Nuculidae, Solemyidae, and Nuculanidae, representative live images **A** *Platidia anomioides* sp. inc. (B211, exterior view) **B** *Platidia anomioides* sp. inc. (B211, interior view) **C** *Ennucula colombiana* (M17879) **D** *Ennucula* stet. (M16966) **E** *Nucula chrysocoma* (M11989) **F** *Acharax* cf. *johnsoni* (M15768, adult specimen, opened) **G** *Acharax* cf. *johnsoni* (M17077, young specimen) **H** *Nuculana* cf. *callimene* (M17063). Scale bars: 1 mm (**A–E, H**); 1 cm (**F, G**).

(2006). M17007 also grouped with the “*Acharax* 1” clade of (Fukasawa et al. 2017), showing 99.61–99.94% identity to sequences from the Aleutian Trench ([AJ563760.1](#)), Java Trench ([AJ563756.1](#), [AJ563757.1](#)), Chishima Trench seeps ([LC186962.1](#)), Japan Trench seeps ([LC186965.1](#), [LC186966.1](#)), Hine Hina vents in the Lau Basin ([LC186970.1](#)), and Haima seeps ([OQ836650.1](#)). M15768 from

Mound 12 (~ 1000 m) grouped with the “*Acharax 2*” clade of Fukasawa et al. (2017), showing 99.14–99.77% identity to sequences from seeps off Oregon ([AJ563751.1](#), [AJ563752.1](#), [AJ563753.1](#), [AJ563754.1](#), [AJ563755.1](#)), off Peru ([AJ563762.1](#)), and at the Nankai Trough ([LC186957.1](#)). The “*Acharax 1*” and “*Acharax 2*” clades show the same depth range globally (765–5300 m and 780–5300 m, respectively) (Fukasawa et al. 2017), so further sequencing and sampling are needed to assess depth stratification of *Acharax* at the CRM. A detailed comparison of these clades may also illuminate distinguishing morphological features.

CRM specimens from Mound 11 and Mound 12 are hosts of copepods (see *Cyclopoida* sp. SIO_BIC_C12780), the chrysopetalid *Natsushima sashai* (Aguado and Rouse 2011), and the hesionid *Neogyptis jeffruocoi* (Rouse et al. 2018).

Mollusca | Bivalvia | Protobranchia | Nuculanida | Nuculanidae

***Nuculana* cf. *callimene* (Dall, 1908)**

Fig. 27H

Material examined. S0220: M17063.

Localities. Subduction Plume (3434 m).

Remarks. This specimen is morphologically similar to *N. callimene* except for the posterior end and may represent an undescribed species. *N. callimene* is known from western Baja California to the Pacific margin of Panama (type locality), 183–3200 m (Coan and Valentich-Scott 2012; Hendrickx et al. 2016).

***Nuculana* cf. *hamata* (Carpenter, 1864)**

Fig. 28A

Material examined. AD4510: M12047.

Localities. Jacó Summit (742 m).

Remarks. This specimen resembles *N. hamata*, which is known from Alaska to the Gulf of California (type locality: Catalina Island, southern California), 20–1100 m, and may include cryptic species; records of this morphologically variable species from further south are considered suspect and warrant further investigation (Coan and Valentich-Scott 2012; Hendrickx et al. 2016). Genetic comparison to *N. cf. hamata* recorded from Baja California, 750–850 m (Hendrickx et al. 2016), may be informative.

Mollusca | Bivalvia | Protobranchia | Nuculanida | Bathyspinulidae

***Tindariopsis grasslei* (Allen, 1993)**

Fig. 28B

Material examined. AD4503: M11985 (dead shell only); AD4506: M12005; AD4590: M12141; AD4971: M16735, M16736; AD4973: M16749, M16750; AD4977: M16806; AD4987: M16902; AD4985: M16908; AD4988: M16964; S0217: M17042.

Localities. Mound 12 (990 m, dead shell; 992–1010 m, live specimens), Mound 11 (1009 m), Parrita Seep (~ 1030–1179 m), The Thumb (1069 m), Jacó Scar (1783–1817 m).

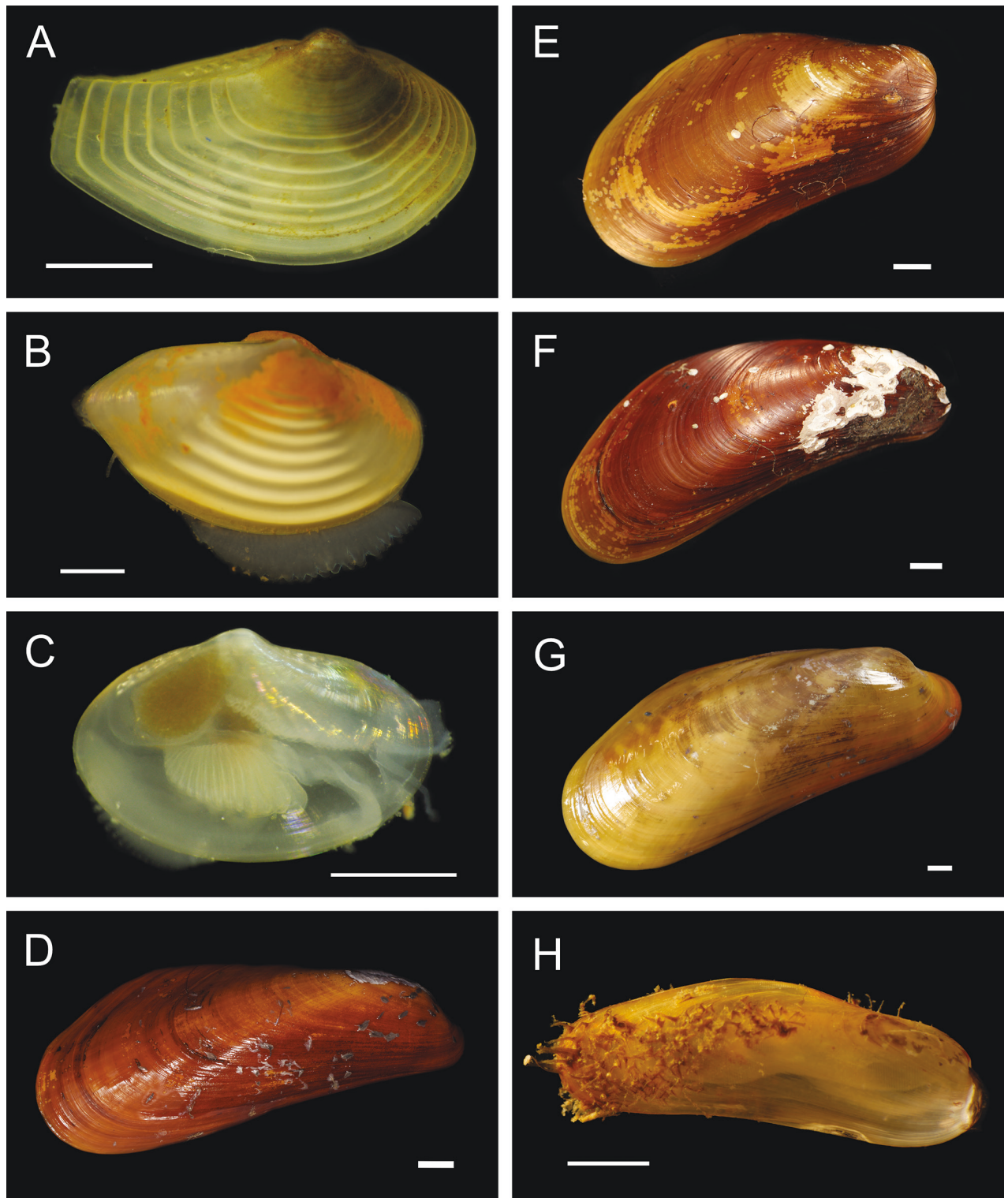


Figure 28. Mollusca: Bivalvia: Nuculanidae, Bathyspinulidae, Yoldiidae, and Mytilidae, representative live images **A** *Nuculana* cf. *hamata* (M12047) **B** *Tindariopsis grasslei* (M16806) **C** *Yoldiella* stet. (M12046) **D** *Bathymodiolus billschneideri* (M12074) **E** *Bathymodiolus earlougheri* (M14479) **F** *Bathymodiolus nancyschneiderae* (M14531) **G** *Bathymodiolus thermophilus* (M17023) **H** *Idas* stet. (M17069). Scale bars: 1 mm (**A–C, H**); 1 cm (**D–G**).

Distribution. Known from the Guaymas Basin, Gulf of California (type locality, 2003 m), to the Costa Rica Subduction Zone, 1400–2012 m (Coan and Valentinich-Scott 2012).

New records. CRM specimen M16908 from 992 m represents a new minimum depth for this species.

Remarks. This species is placed “with reluctance” in *Tindariopsis* (Coan and Valentich-Scott 2012). Preliminary molecular work suggests a position closer to *Malletia*. M11985 was associated with a naturally occurring wood fall.

Mollusca | Bivalvia | Protobranchia | Nuculanida | Yoldiidae

Yoldiella stet.

Fig. 28C

Material examined. AD4510: M12046.

Localities. Jacó Summit (742 m).

Mollusca | Bivalvia | Pteriomorphia | Mytilida | Mytilidae

***Bathymodiolus billschneideri* McCowin, Feehery & Rouse, 2020**

Fig. 28D

Reference. McCowin et al. 2020**.

Localities. Parrita Seep (~ 1400 m), Jacó Scar (~ 1750–1900 m; type locality).

Distribution. Known only from the CRM seeps and apparently found no shallower than ~ 1400 m.

Remarks. *B. billschneideri* is a host of the scaleworms *Branchiopolynoe eliseae*, *Br. halliseyae*, *Br. kajsae*, and *Br. meridae* (Lindgren et al. 2019). Previous reports of mytilid mussels at Parrita Seep (previously published as “Quepos Seep”) (Sahling et al. 2008) and brown-colored *Bathymodiolus* mussels at Jacó Scar (Levin et al. 2012, 2015) are now known to correspond to *B. billschneideri*.

***Bathymodiolus earlougheri* McCowin, Feehery & Rouse, 2020**

Fig. 28E

Reference. McCowin et al. 2020**.

Localities. Mound 12 (~ 1000 m), The Thumb (1073 m), Jacó Scar (~ 1750–1900 m; type locality).

Distribution. Known only from the CRM seeps.

Remarks. *B. earlougheri* likely also occurs at sites of intermediate depth (e.g., Parrita Seep, ~ 1400 m) but has not been collected there, likely due to limited sampling (McCowin et al. 2020). *B. earlougheri* is the only known host of the chrysopetalid *Laubierus alvini* (Aguado and Rouse 2011). It is also a host of the scaleworms *Branchiopolynoe halliseyae*, *Br. kajsae*, *Br. meridae*, and possibly *Br. eliseae* (Lindgren et al. 2019). Previous reports of golden-colored *Bathymodiolus* mussels at multiple CRM seep sites (Levin et al. 2012, 2015) are now known to correspond to *B. earlougheri*.

***Bathymodiolus nancyschneiderae* McCowin, Feehery & Rouse, 2020**

Fig. 28F

Reference. McCowin et al. 2020**.

Localities. Mound 12 (~ 1000 m), Jacó Slope (1063 m; type locality), The Thumb (1073 m).

Distribution. Known only from the CRM seeps and apparently found no deeper than ~ 1000 m.

Remarks. *B. nancyschneiderae* is a host of the scaleworms *Branchipolynoe eliseae*, *Br. halliseyae*, *Br. kajsae*, and possibly also *Br. meridae* (Lindgren et al. 2019). Previous reports of brown-colored *Bathymodiolus* spp. at Mound 11 and Mound 12 (Levin et al. 2015) are now known to correspond to *B. nancyschneiderae*.

***Bathymodiolus thermophilus* Kenk & B. R. Wilson, 1985**

Fig. 28G

References. Levin et al. 2012; McCowin et al. 2020.

Localities. Jacó Scar (1794–1814 m).

Distribution. Originally described from 2495 m at the Galápagos Rift (Kenk and Wilson 1985) and distributed along the East Pacific Rise between 13°N and 21°S latitude (Johnson et al. 2013), *B. thermophilus* was previously thought to be restricted to hydrothermal vents but also occurs at the Jacó Scar “hydrothermal seep” site (Levin et al. 2012; McCowin et al. 2020).

Remarks. *B. thermophilus* specimens from the CRM seeps were not observed to contain *Branchipolynoe* spp. scaleworms, but this apparent absence may reflect limited sample size and the tendency of the worms to evacuate the mussels upon disturbance.

Idas stet.

Fig. 28H

Reference. Xu et al. (2017) for phylogenetic analysis.

Material examined. AD4587: M13006 ([KU975037](#)); AD4907: M16100; S0219: M17069 ([PQ450399](#)).

Localities. Rio Bongo Scar (661 m; this study), Mound 12 (996–999 m).

Remarks. Associated with naturally occurring and experimentally deployed wood. As described in Xu et al. (2017), M13006 shows morphological similarity to *Idas washingtonius* (Bernard, 1978), which occurs in the eastern Pacific from Washington to the Guaymas Basin and in the western Pacific, 1240–2200 m (Coan et al. 2000), but genetically it is more closely related to *I. macdonaldi* Gustafson, Turner, Lutz & Vrijenhoek, 1998, which occurs at hydrocarbon seeps in the Gulf of Mexico (Gustafson et al. 1998). The paraphyly of *Idas* (Lorion et al. 2013; Xu et al. 2017) warrants further investigation.

Mollusca | Bivalvia | Pteriomorphia | Pectinida | Pectinidae

***Delectopecten vancouverensis* (Whiteaves, 1893)**

Fig. 29A

Material examined. AD4512: M12063; AD4918: M16184.

Localities. Quepos Slide (338–~ 400 m).

Distribution. Originally described from British Columbia, Canada; recorded in the eastern Pacific from Alaska to Baja California as well as the Bering Sea and northwestern Pacific to the Sea of Japan, 20–4100 m (Coan and Valentich-Scott 2012; Kamenev 2013; Hendrickx et al. 2016). GBIF includes a museum record from the Galápagos Islands, 466 m (Blum and Fong 2016c).

New records. Pending verification of the GBIF record, the CRM specimens represent new southern records for this species.

Remarks. M16184 was associated with epibiotic hydroids. DNA sequences could not be obtained.

***Delectopecten zaca* (Hertlein, 1935)**

Fig. 29B

Material examined. AD4921: M16188.

Localities. Quepos Slide (~ 345–394 m).

Distribution. Known from Baja California (type locality: Cabo San Lucas, 37–402 m) (Hertlein 1935), to Peru and the Galápagos Islands, 10–1840 m (Coan and Valentich-Scott 2012).

Remarks. DNA sequences could not be obtained.

Mollusca | Bivalvia | Heterodonta | Lucinida | Thyasiridae

***Thyasira methanophila* P. G. Oliver & Sellanes, 2005**

Fig. 29C

Reference. Coan and Valentich-Scott (2012), in which the locality of specimen SBMNH 350533 (9.033, -84.621; 1408 m; no DNA sequences; Santa Barbara Museum of Natural History) aligns with the locality known in this work as Parrita Seep.

Additional material examined. S0230: M17098 (juvenile specimen, only valves collected).

Localities. Parrita Seep (1408 m); Mound Jaguar (1909 m; this study, valves).

Distribution. Also known from seeps at the type locality off Concepción, central Chile, 780 m (Oliver and Sellanes 2005). Confirmation of live specimens at Mound Jaguar would establish a new depth record.

Thyasira stet.

Fig. 29D

Material examined. AD4512: M12059 ([PQ450412](#)); AD4978: M16840; AD4988: M16977.

Localities. Quepos Slide (~ 344–411 m), Mound 12 (~ 996–999 m), Jacó Scar (1783 m).

Remarks. This morphospecies warrants comparison to other eastern Pacific specimens that have been dubiously reported as *Thyasira flexuosa* (Montagu, 1803) (type locality: Great Britain) and likely represent several cryptic species (Coan et al. 2000; Coan and Valentich-Scott 2012). For M12059, the closest COI BLASTN results on GenBank were *Thyasira sarsii* (R. A. Philippi, 1845) from Norway ([AM706508.1](#) and [AM706509.1](#), both 97.09% identity). Based on a recent phylogeny (Fukasawa et al. 2017), *Thyasira* appears to require taxonomic revision.

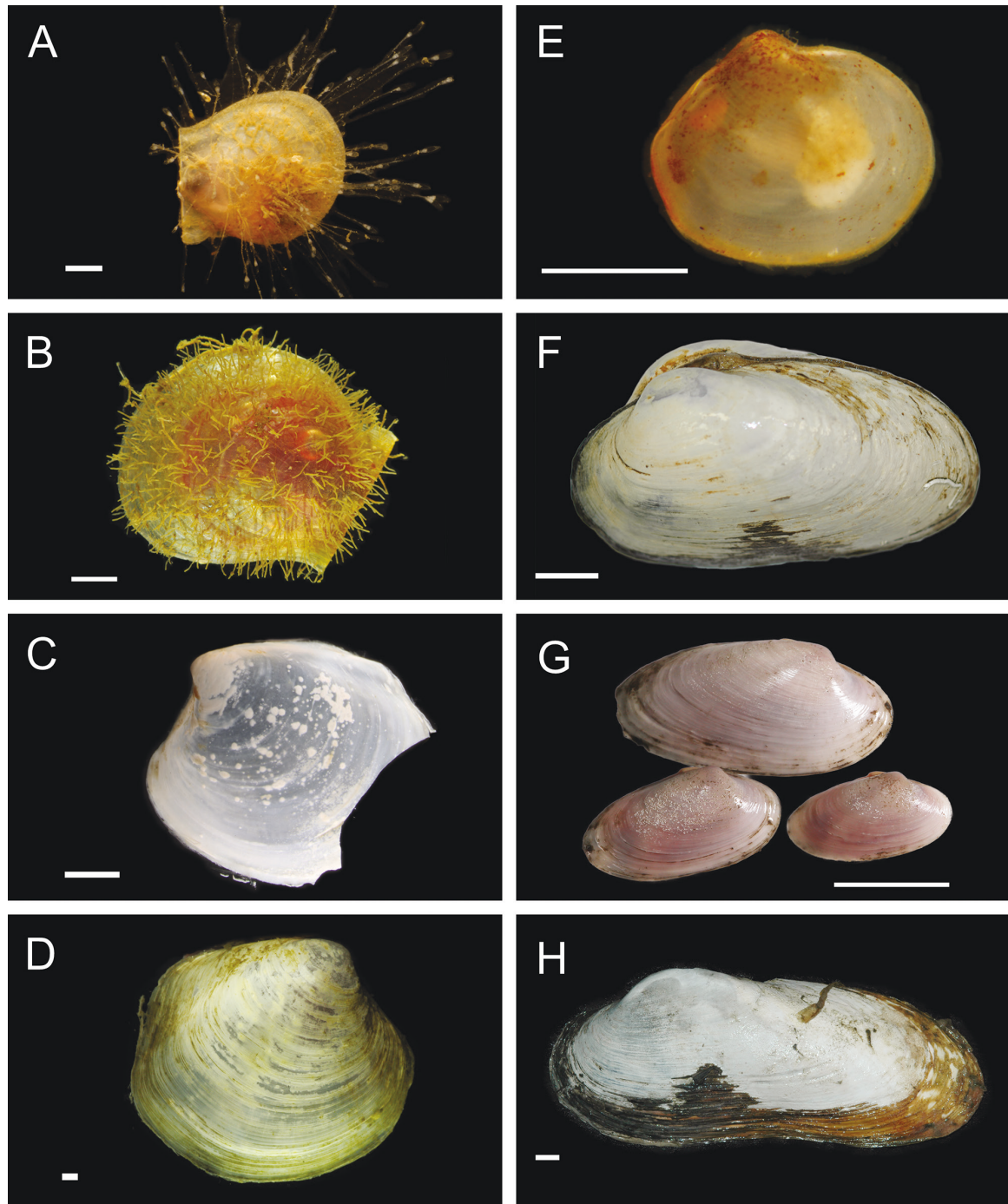


Figure 29. Mollusca: Bivalvia: Pectinidae, Thyasiridae, Galeommatidae, and Vesicomysidae, representative live images **A** *Delectopecten vancouverensis* (M16184) **B** *Delectopecten zacae* (M16188) **C** *Thyasira methanophila* (M17098, valve only) **D** *Thyasira stet.* (M12059) **E** *Axinodon cf. redondoensis* (M16845) **F** *Archivesica gigas* (M12011) **G** *Archivesica* sp. 6 sec. Audzi-jonyte et al. 2012 (M16141) **H** *Archivesica* sp. SIO_BIC_M12070 aff. *gigas* (M12070). Scale bars: 1 mm (**A–E**); 1 cm (**F–H**).

Mollusca | Bivalvia | Heterodonta | Galeommatida | Galeommatidae

***Axinodon cf. redondoensis* (T. A. Burch, 1941)**

Fig. 29E

Material examined. AD4978: M16845.

Localities. Mound 12 (~ 996–999 m).

Remarks. Likely an undescribed species. Possibly a range and depth extension of *Axinodon redondoensis*, presently known from Washington to Redondo Beach, southern California (type locality, 137 m), 120–330 m (Harry 1969; Valentich-Scott 1998; Coan et al. 2000).

Mollusca | Bivalvia | Heterodonta | Venerida | Vesicomysidae

We thank Elena Krylova (P.P. Shirshov Institute of Oceanology, Russian Academy of Sciences) for input on this section.

***Archivesica gigas* (Dall, 1896)**

Fig. 29F

References. Audzijonyte et al. 2012; Levin et al. 2012.

Additional material examined. AD4506: M12007; AD4507: M12011 ([PQ449002](#)); AD4508: M12014 ([PQ449401](#)).

Localities. Parrita Seep (1186 m, ~ 1400 m; this study), Parrita Scar (~ 1659–1667 m; this study), Jacó Scar (~ 1800 m); an unnamed locality ~ 85 km northwest of Mound Jaguar (10.3000, -86.3053; 1531 m) (Audzijonyte et al. 2012).

Distribution. Originally described from the Guaymas Basin, Gulf of California, 1567 m (Dall 1896) and reported as far north as the Gulf of Alaska (Coan and Valentich-Scott 2012). Genetically confirmed records are known from seeps, sedimented vents, and whale falls from the Oregon Subduction Zone to Costa Rica, 1013–2028 m, as well as a seep at 1200 m in Hiroo Submarine Canyon off Hokkaido (Audzijonyte et al. 2012).

***Archivesica* sp. 6 sec. Audzijonyte et al. 2012**

Fig. 29G

Reference. Audzijonyte et al. (2012) for DNA sequences and phylogenetic analysis.

Additional material examined. AD4590: M13509 ([PQ450381](#), [PQ450398](#); tissues); AD4912: M16141 ([PQ449413](#)).

Localities. Jacó Scar (1677 m; ~ 1791–1842 m, this study).

Distribution. Known only from the CRM seeps.

Remarks. Morphologically resembles *A. gigas* and may warrant description as a new species (Audzijonyte et al. 2012).

***Archivesica* sp. 7 sec. Audzijonyte et al. 2012**

References. Peek et al. (2000) and Audzijonyte et al. (2012) for DNA sequences and phylogenetic analysis. Not collected in this study.

Localities. An unnamed seep at 3002 m (9.69850, -86.06817) and a “low-temperature vent” at 3096 m (9.7112, -86.0777) (Audzijonyte et al. 2012), both ~ 20 km west of Mound Jaguar.

Distribution. Also known from a seep at 4300 m in the Middle America Trench off Mexico (Audzijonyte et al. 2012; Decker et al. 2012) and from the Peru margin, 3711–4724 m (Audzijonyte et al. 2012).

Remarks. This taxon warrants consideration as a new species based on COI sequences, but no morphological voucher specimens are available (Audzijonyte et al. 2012).

***Archivesica* sp. SIO_BIC_M12070 aff. *gigas* (Dall, 1896)**

Fig. 29H

Material examined. AD4513: M12070 ([PQ450402](#)).

Localities. Jacó Scar (1744 m).

Remarks. An undescribed species. The COI sequence of this specimen was 94.83–95.20% identical to sequences of *A. gigas* from northern Japan ([PP899629.1](#)), the Guaymas Basin, Gulf of California ([MF959623.1](#)) (Liu et al. 2018), [MT947383.1](#)), and off northern California ([MT947382.1](#)).

***Calyptogena costaricana* Krylova & Sahling, 2006**

Reference. Krylova and Sahling 2006**. Not collected in this study.

Localities. Mound 10 (2258–2263 m; type locality).

Distribution. Also known from seeps in Monterey Canyon, off California, 2193–2219 m, and a vent on the Peru margin, 2500 m (Audzijonyte et al. 2012).

***Calyptogena diagonalis* Barry & Kochevar, 1999**

Fig. 30A

References. Barry and Kochevar 1999**; Audzijonyte et al. 2012.

Additional material examined. S0220: M17064, M17065.

Localities. Subduction Plume (3408 m; this study); an unnamed seep (9°42.28'N, 86°4.38'W; 2980–3800 m; type locality) (Barry and Kochevar 1999) and a “low-temperature vent” (9.7112, -86.0777; 3096 m) (Audzijonyte et al. 2012), both ~ 20 km west of Mound Jaguar.

Distribution. Also known from seeps at the Oregon margin, 2021–2089 m (Barry and Kochevar 1999; Audzijonyte et al. 2012; Coan and Valentich-Scott 2012), vents at the Juan de Fuca Ridge, 2400–2437 m (Peek et al. 1997; Audzijonyte et al. 2012), and the Peru margin, 2290–3070 m (Audzijonyte et al. 2012).

Remarks. M17065 was associated with an anemone, Co3084. Based on morphological and genetic similarities, Audzijonyte et al. (2012) recommend synonymization of this species (referenced as *Archivesica diagonalis*) with the western Pacific nominal species *A. magnocultellus* (Okutani, Kojima & Iwasaki, 2002), pending examination of the type material of *A. magnocultellus*. Formal synonymization would extend the species' range to seeps off southern Japan, 1900–2500 m (Audzijonyte et al. 2012).

***Calyptogena* sp. 3 sec. Audzijonyte et al. 2012**

Reference. Audzijonyte et al. (2012) for DNA sequences and phylogenetic analysis. Not collected in this study.

Localities. Unnamed seep ~ 25 km west of Mound Jaguar (9.6678, -86.1193; 3560 m) (Audzijonyte et al. 2012).

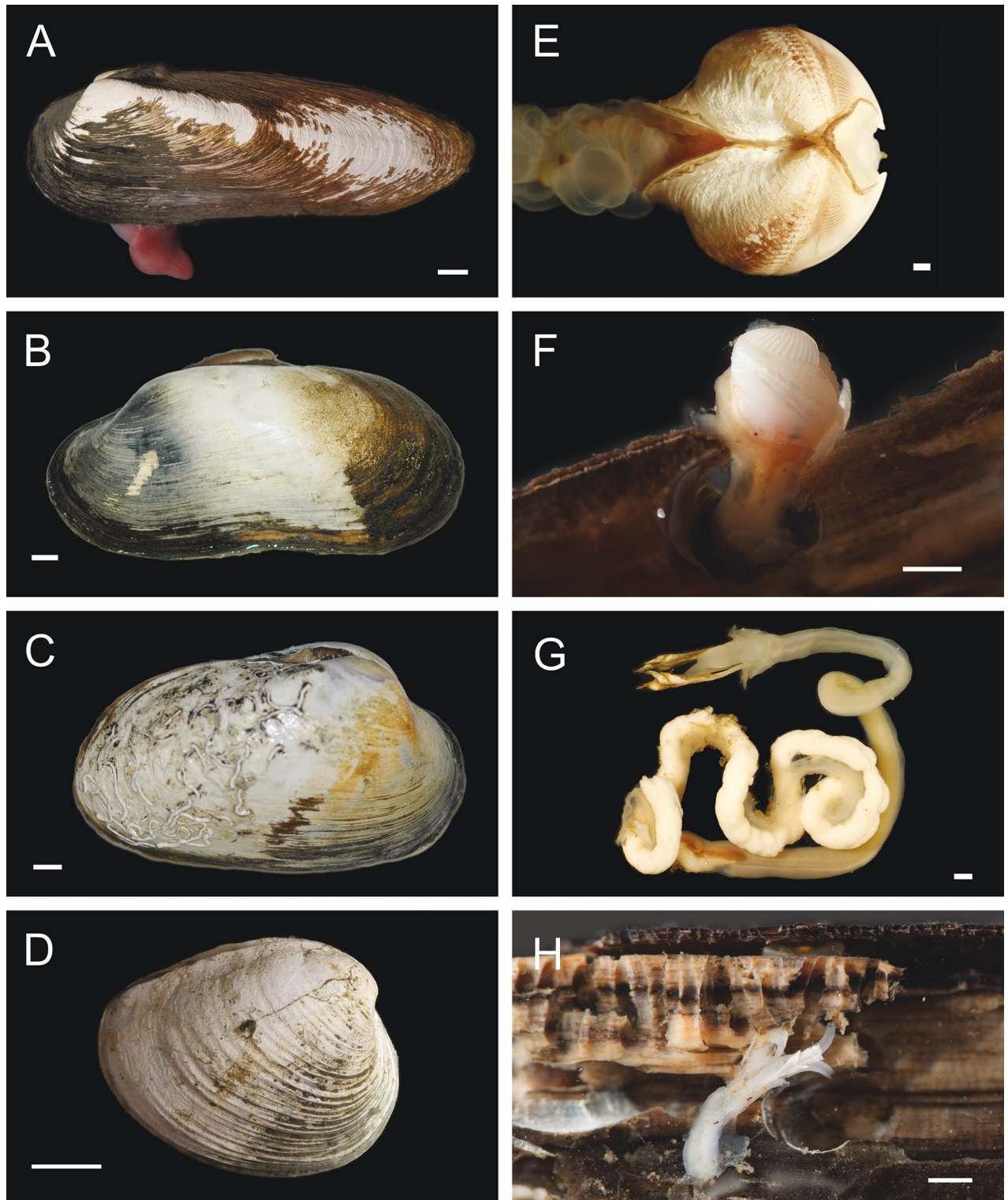


Figure 30. Mollusca: Bivalvia: Vesicomyidae, Xylophagidae, and Teredinidae, representative live images **A** *Calyptogena diagonalis* (M17064) **B** *Phreagena soyoae* (M11999) **C** *Phreagena* sp. 5 sec. Audzijonyte et al. 2012 (M12009) **D** *Pliocardia krylovata* (M17026) **E** *Xylophaga stet.* (M16099) **F** *Xyloreda* gen. inc. (M17870) **G** *Teredinidae* sp. SIO_BIC_M17100 (M17100) **H** *Teredinidae* stet. (M12290). Scale bars: 1 cm (**A–D**); 1 mm (**E–H**).

Distribution. Also known from seep and whale fall habitats off central California, 2895–3449 m (Audzijonyte et al. 2012).

Remarks. This taxon warrants consideration as a new species based on COI sequences, but no morphological voucher specimens are available (Audzijonyte et al. 2012).

***Calyptogena* sp. mt-V sec. Goffredi et al. 2003**

References. Goffredi et al. (2003) and Audzijonyte et al. (2012) for DNA sequences and phylogenetic analysis. Not collected in this study.

Localities. Unnamed “low-temperature vent” ~ 20 km west of Mound Jaguar (9.7112, -86.0777; 3096 m) (Audzijonyte et al. 2012).

Remarks. The single genetic sample of this taxon was originally identified as *Calyptogena pacifica* (Peek et al. 2000) and underwent several nomenclatural revisions upon further phylogenetic analyses (Goffredi et al. 2003; Audzijonyte et al. 2012). This taxon warrants consideration as a new species based on COI sequences, but no morphological voucher specimens are available (Audzijonyte et al. 2012).

***Phreagena extenta* (Krylova & Moskalev, 1996)**

Reference. Peek et al. (2000) for DNA sequences and phylogenetic analysis. Not collected in this study.

Localities. Unnamed seep ~ 20 km west of Mound Jaguar (9.6983, -86.0682; 3002 m).

Distribution. Originally described from a seep in Monterey Bay at 3041 m; recorded from the Gulf of Alaska to Costa Rica, 2889–4445 m (Peek et al. 2000; Audzijonyte et al. 2012; Coan and Valentich-Scott 2012), as well as the Kuril Trench, 3512 m (Okutani et al. 2009; Audzijonyte et al. 2012).

Remarks. Previously placed in *Ectenagena* and then *Calyptogena* (Coan and Valentich-Scott 2012), this species is provisionally, and with an acknowledgment of doubt, allocated to *Phreagena* (Johnson et al. 2017).

***Phreagena soyoae* (Okutani, 1957)**

Fig. 30B

Reference. Audzijonyte et al. (2012) for DNA sequences and phylogenetic analysis.

Additional material examined. AD4505: M11999 ([PQ449397](#), [PQ449398](#), [PQ449399](#), [PQ450407](#), [PQ450408](#)).

Localities. Mound 11 (~ 1019–1025 m; this study); Mound Carablanca, Nicaragua margin (1432 m).

Distribution. Originally described from seeps off Japan, 750–1500 m (type locality: Sagami Bay, 750 m), *P. soyoae* is found at seep, vent, and whale fall habitats from the Juan de Fuca Ridge to the Costa Rica and Nicaragua margins, 519–2400 m (Audzijonyte et al. 2012).

***Phreagena* sp. 5 sec. Audzijonyte et al. 2012**

Fig. 30C

Reference. Audzijonyte et al. (2012) for DNA sequences and phylogenetic analysis.

Additional material examined. AD4506: M12009 ([PQ449400](#)), M12085 ([PQ450380](#)); AD4508: M12088 ([PQ450379](#)); AD4988: M16960 ([PQ449417](#)).

Localities. Mound 11 (1012 m; this study), Jacó Slope (1024 m), Parrita Seep (1186 m and ~ 1400 m, this study; 1408 m). The locality “Mound Quepos” at 1408 m and the upslope region of Jacó Scar at 1024 m in Audzijonyte et al. (2012) are reported as “Parrita Seep” and “Jacó Slope,” respectively, in this work.

Remarks. Morphologically resembles *Phreagena soyoae* and may warrant description as a new species (Audzijonyte et al. 2012).

***Pliocardia krylovata* A. M. Martin & Goffredi, 2012**

Fig. 30D

Reference. Martin and Goffredi 2012**.

Additional material examined. S0213: M17026.

Localities. Jacó Summit (~ 741–744 m; type locality), Mound 12 (~ 967–995 m).

Distribution. Known only from the CRM seeps.

Remarks. Found in sediments near microbial mats and typically positioned with approximately half the shell length above the sediment, forming “a spatial bridge between the oxic overlying water and the sulphide-rich sediment” (Martin and Goffredi 2012). Assignment of this species to the genus *Pliocardia* was tentative based on uncertainty within Pliocardiinae (Martin and Goffredi 2012), and the clade containing this taxon was later suggested to warrant a new genus (Johnson et al. 2017).

Mollusca | Bivalvia | Heterodonta | Myida | Pholadoidea | Xylophagaidae

***Xylophaga* stet.**

Fig. 30E

Material examined. AD4906: M15767, M16099, M16111.

Localities. Mound 12 (1002 m).

Remarks. Associated with experimentally deployed wood. We thank Chiara Romano (University of Gastronomic Sciences) for this identification.

***Xyloredo* gen. inc.**

Fig. 30F

Material examined. AD4588: M17870.

Localities. Mound 12 (~ 1000 m).

Remarks. Associated with a naturally occurring palm wood fall.

Mollusca | Bivalvia | Heterodonta | Myida | Pholadoidea | Teredinidae

We thank Reuben Shipway (University of Plymouth) for these identifications.

Teredinidae sp. SIO_BIC_M17100

Fig. 30G

Material examined. AD4913: M16147; S0230: M17100 ([PQ449424](#)).

Localities. Jacó Scar (1817 m), Mound Jaguar (1896 m).

Remarks. An undescribed species associated with naturally occurring wood falls.

Teredinidae stet.

Fig. 30H

Material examined. AD4588: M12290.

Localities. Mound 12 (995 m).

Remarks. Associated with a naturally occurring wood fall. DNA sequences could not be obtained.

Mollusca | Bivalvia | Heterodonta | Anomalodesmata | Cuspidariidae

***Bathyneera tillamookensis* (Dall, 1916)**

Fig. 31A

Material examined. AD4989: M16982.

Localities. Jacó Scar (1762 m).

Distribution. Originally described from the Oregon margin, 1438 m, and known from British Columbia, Canada, to Peru as well as the Atlantic and New Zealand, 439–2850 m (Coan and Valentich-Scott 2012).

***Luzonia chilensis* (Dall, 1890)**

Fig. 31B

Material examined. AD4508: M12026; AD4974: M16769; AD4977: M16808; AD4989: M16980, M16981, M16983.

Localities. Mound 12 (990m), Parrita Seep (1402m), Jacó Scar (1762–1783m).

Distribution. Originally described from Isla Mocha, central Chile, 1238 m, and known from the U.S. Pacific coast off Washington to southern Chile, 100–1875 m (Coan and Valentich-Scott 2012).

Mollusca | Gastropoda

We list the six widely accepted major gastropod clades (Bouchet et al. 2017) in a sequence reflecting their current phylogenomic relationships (Uribe et al. 2022; Zhong et al. 2022), acknowledging the possibility of further revision of higher-level clade names. Within these clades we organize the listings following WoRMS. We recognize the need for considerable further taxonomic and genetic work, especially on limpet-shaped forms such as Neolepetopsidae and Cocculiniformia (Desbruyères et al. 2006; Chen et al. 2019b, 2023). Several gastropod morphospecies occurring at Jacó Scar were provisionally identified in Electronic supplemental table S6 of Levin et al. (2012) and are here linked to voucher specimens where possible.

Mollusca | Gastropoda | Patellogastropoda | Lottioidea | Lepetidae

***Iothia* stet.**

Fig. 31C, D

Reference. Electronic supplemental table S6 of Levin et al. (2012) (occurrences only, Jacó Scar).

Material examined. AD4508: M12025 ([PQ450378](#)); AD4916: M16172, M16173.

Localities. Parrita Seep (~ 1401–1419 m), Jacó Scar (1611 m).

Remarks. For M12025, the closest COI BLASTN results on GenBank were specimens of *Lothia fulva* (O. F. Müller, 1776) from the North Sea ([KR084663.1](#), [KR084581.1](#); 94.53% identity).

Mollusca | Gastropoda | Patellogastropoda | Lottioidea | Neolepetopsidae

***Eulepetopsis* gen. inc.**

Fig. 31E, F

Reference. Electronic supplemental table S6 of Levin et al. (2012) as *Eulepetopsis* sp. (occurrences only, Jacó Scar).

Material examined. AD4506: M12004; AD4590: M17882, M18921, M19153.

Localities. Parrita Seep (1186 m), Jacó Scar (~ 1800 m).

Remarks. Morphologically similar to *Eulepetopsis*, although the two described species of *Eulepetopsis* are known only from high-temperature vents and Neolepetopsidae may require revision (Chen et al. 2023).

Neolepetopsidae stet.

Fig. 31G, H

Reference. Several morphospecies were reported in Electronic supplemental table S6 of Levin et al. (2012) (occurrences only, Jacó Scar).

Material examined. AD4501: M11963; AD4502: M12093; AD4586: M18777, M18780, M18800, M18802, M18804, M18807; AD4587: M19015, M19116, M19155, M19156; AD4588: M19019, M19020, M19025, M19031, M19034, M19035, M19040, M19041, M19060, M19065, M19071, M19097, M19108, M19138, M19046, M19050, M19053, M19064, M19070, M19084, M19154; AD4589: M18813, M18827, M18839, M18842, M18848, M18858, M18864, M18910, M18918, M18926, M18929, M18958, M18964, M18966, M19139, M19140, M19141, M19157; AD4590: M18990; AD4591: M18874, M18882, M18885, M18892, M18902, M18903, M19163.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. Under genetic and morphological investigation (Betters et al. (2024), in press at the time of this work's acceptance). Neolepetopsidae and *Paralepetopsis* appear to be paraphyletic (Chen et al. 2023).

***Paralepetopsis* cf. *clementensis* J. H. McLean, 2008**

Fig. 32A

Reference. Reported in Electronic supplemental table S6 of Levin et al. (2012) as "*Neolepetopsis* [sic] *clementensis*" (occurrences only).

Material examined. AD4590: M18930, M18935, M18937, M18961, M18969, M18976, M18980, M18989, M19142; AD4591: M18884, M18901.

Localities. Jacó Scar (~ 1800 m).

Distribution. *Paralepetopsis clementensis* is known only from a whale fall off southern California at 1800 m depth (McLean 2008). Genetic confirmation of the CRM seep occurrences would represent new southern records and the

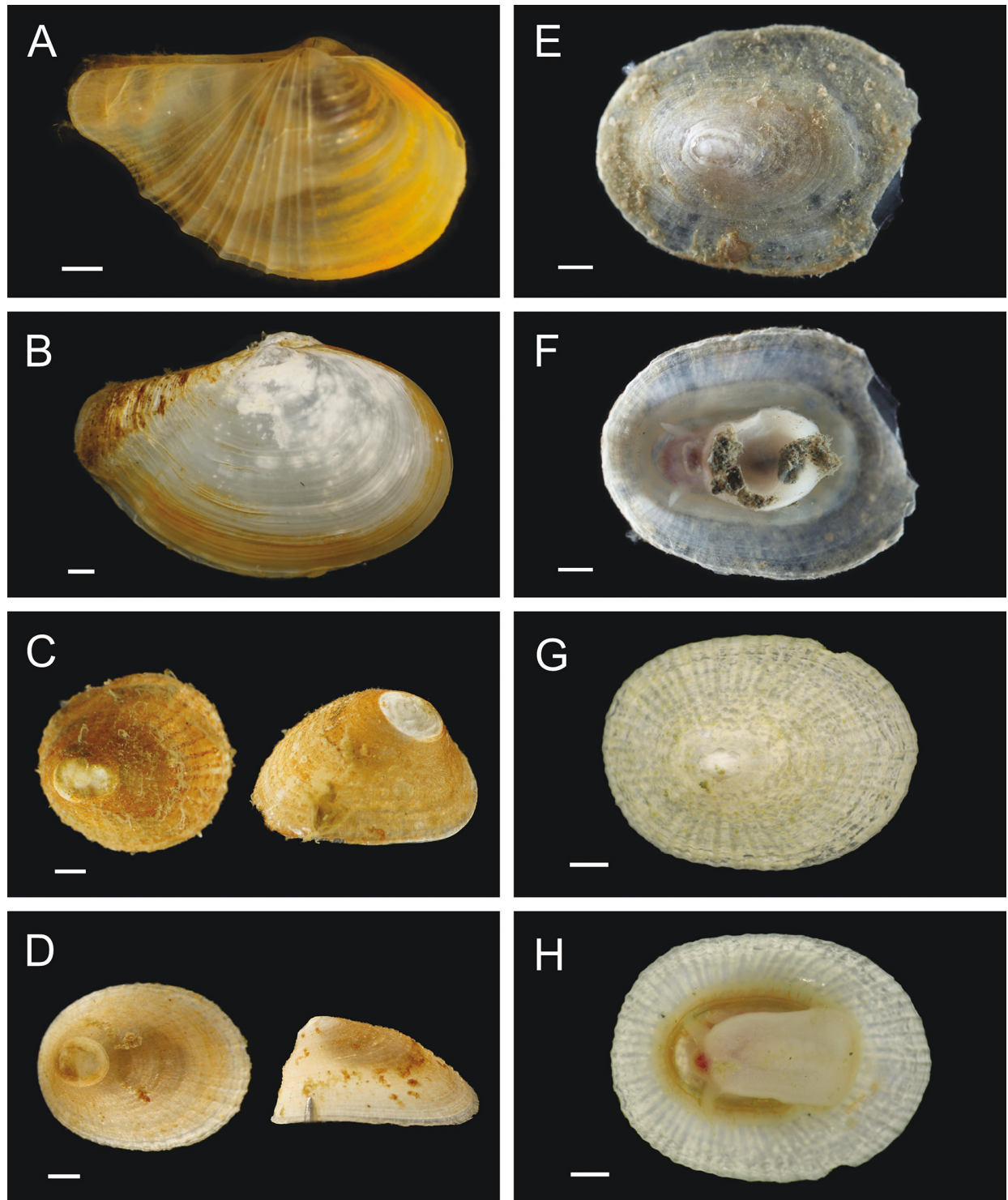


Figure 31. Mollusca: Bivalvia and Gastropoda: Patellogastropoda, representative live images **A** *Bathyneaea tillamookensis* (M16982) **B** *Luzonia chilensis* (M16808) **C** *Iothia* stet. (M16172, separate specimens in dorsal and lateral view) **D** *Iothia* stet. (M16173, separate specimens in dorsal and lateral view) **E** *Eulepetopsis* gen. inc. (M17882, dorsal view) **F** *Eulepetopsis* gen. inc. (M17882, ventral view) **G** *Neolepetopsidae* stet. (M11963, dorsal view) **H** *Neolepetopsidae* stet. (M11963, ventral view). Scale bars: 1 mm.

first seep records for this species. Additional specimens of *Paralepetopsis* are under morphological and genetic investigation (Better et al. (2024), in press at the time of this work's acceptance).

Mollusca | Gastropoda | Patellogastropoda | Lottioidea |
Pectinodontidae

***Bathyacmaea* stet.**

Fig. 32B

Reference. Electronic supplemental table S6 of Levin et al. (2012) (occurrences only, Jacó Scar).

Material examined. AD4503: M11987; AD4587: M18812; AD4590: M18925, M18928, M18945, M18953, M18960, M18968, M18978, M18987; AD4591: M18873, M18881, M18888, M18893, M19151.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. Under genetic and morphological investigation (Betters et al. (2024), in press at the time of this work's acceptance).

Mollusca | Gastropoda | Caenogastropoda | Littorinimorpha | Eulimidae

***Fuscapex* stet.**

Fig. 32C

Material examined. AD4506: M12006.

Localities. Parrita Seep (~ 1030–1033 m).

Remarks. The host of this specimen was not determined, but *Fuscapex* is only known as a parasite on ophiuroids (Bouchet and Warén 1986). The specimen was recovered among various animals associated with a coralliid (voucher specimen Co2271), including potential hosts *Ophiacantha moniliformis* (voucher specimen E4390) and *Ophiuroglypha* cf. *meridionalis* (voucher specimen E7978).

Mollusca | Gastropoda | Caenogastropoda | Littorinimorpha | Rissoidae

***Alvania* stet.**

Fig. 32D

Material examined. AD4589: M18834.

Localities. Mound 12 (997 m).

Remarks. Associated with a naturally occurring wood fall.

Mollusca | Gastropoda | Caenogastropoda | Littorinimorpha |
Vitrinellidae

Vitrinellidae stet.

Fig. 32E

Material examined. AD4587: M19091.

Localities. Mound 12 (996 m).

Remarks. Associated with a naturally occurring wood fall.

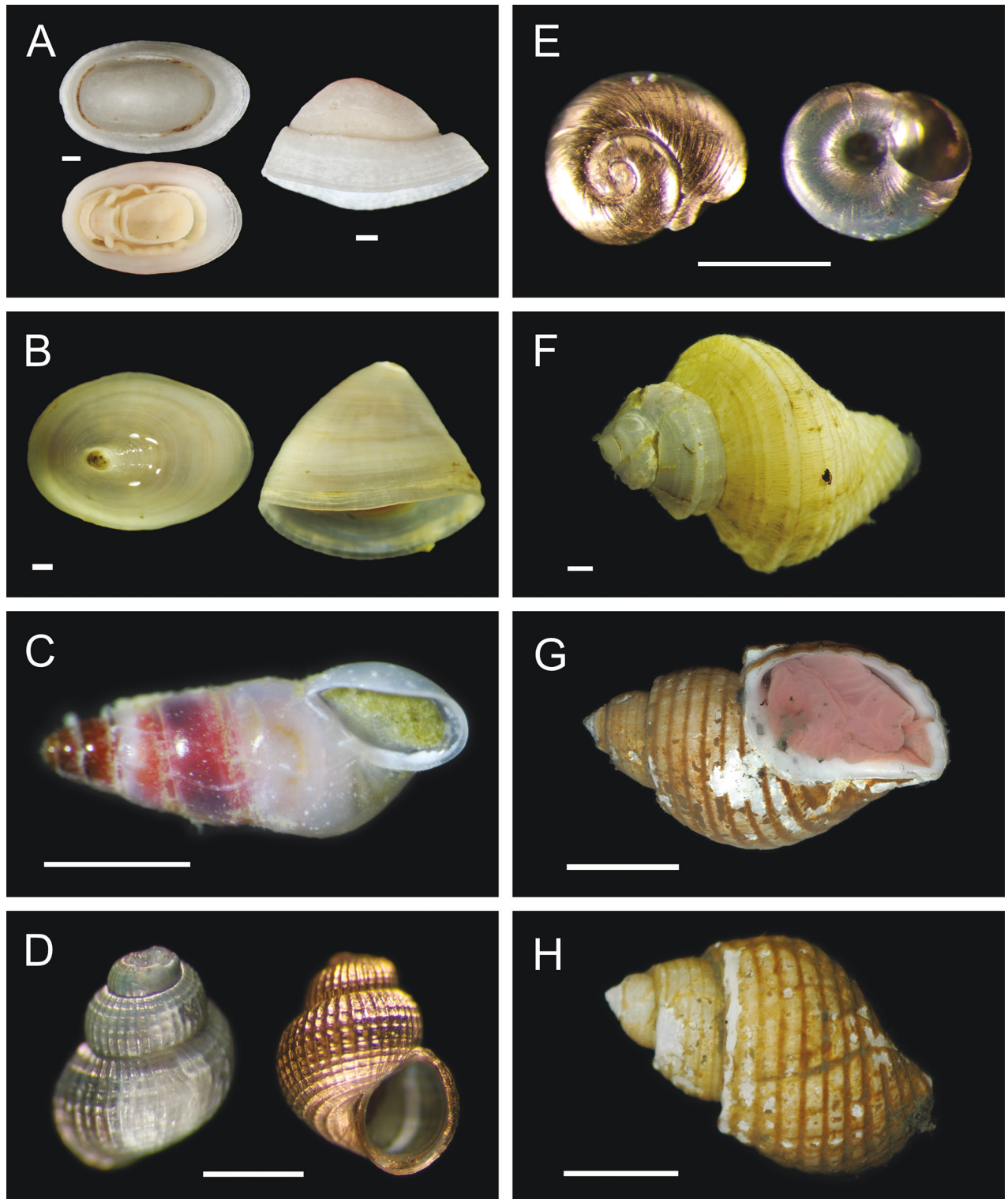


Figure 32. Mollusca: Gastropoda: Patellogastropoda and Caenogastropoda, representative images. Live specimens are depicted unless otherwise specified **A** *Paralepetopsis* cf. *clementensis* (M18937, preserved specimen in dorsal, ventral, and lateral view) **B** *Bathyacmaea* stet. (M11987, same specimen in dorsal and lateral view) **C** *Fuscapex* stet. (M12006) **D** *Alvania* stet. (M18834, same preserved specimen in lateral and apertural view) **E** Vitrinellidae stet. (M19091, same preserved specimen in apical and umbilical view) **F** *Neptunea* stet. (M12023) **G** *Cancellaria* nr. *rosewateri* (M12051, apertural view) **H** *Cancellaria* nr. *rosewateri* (M12051, lateral view). Scale bars: 1 mm (**A–F**); 1 cm (**G, H**).

Mollusca | Gastropoda | Caenogastropoda | Neogastropoda |
Buccinidae

***Neptunea amianta* (Dall, 1890) sp. inc.**

Fig. 32F

Material examined. AD4508: M12023; AD4913: M16143; S0230: M17099.

Localities. Parrita Seep (~ 1401–1419 m), Jacó Scar (1847 m), Mound Jaguar (1908 m).

Remarks. Possibly a southern range extension of *Neptunea amianta*, which was originally described from southern California at 757 m and has been reported from the Bering Sea off Alaska to northern Baja California, 100–3500 m (McLean 1996). Scavenging aggregations of *N. amianta* have been associated with organic falls and seeps (McClain and Nekola 2007).

Mollusca | Gastropoda | Caenogastropoda | Neogastropoda |
Cancellariidae

***Cancellaria* nr. *rosewateri* Petit, 1983**

Fig. 32G, H

Material examined. AD4510: M12051.

Localities. Jacó Summit (744 m).

Remarks. This specimen warrants further comparison to *Cancellaria rosewateri*, described from the Gulf of Mexico, 366 m (Petit 1983).

Mollusca | Gastropoda | Caenogastropoda | Neogastropoda |
Raphitomidae

***Gymnobela* stet.**

Fig. 33A

Material examined. AD4504: M11990 ([PQ449396](#)); AD4988: M16958; S0217: M17041.

Localities. Mound 11 (~ 999–1025 m), The Thumb (1069 m).

Remarks. For M11990, the closest COI BLASTN result on GenBank was a specimen of *Typhlosyrinx* (Raphitomidae) from Papua New Guinea, 680–689 m ([MH308407.1](#), 93.26% identity) (Uribe et al. 2018).

***Phymorhynchus* gen. inc.**

Fig. 33B, C

Material examined. S0214: M17038.

Localities. Jacó Scar (1801 m).

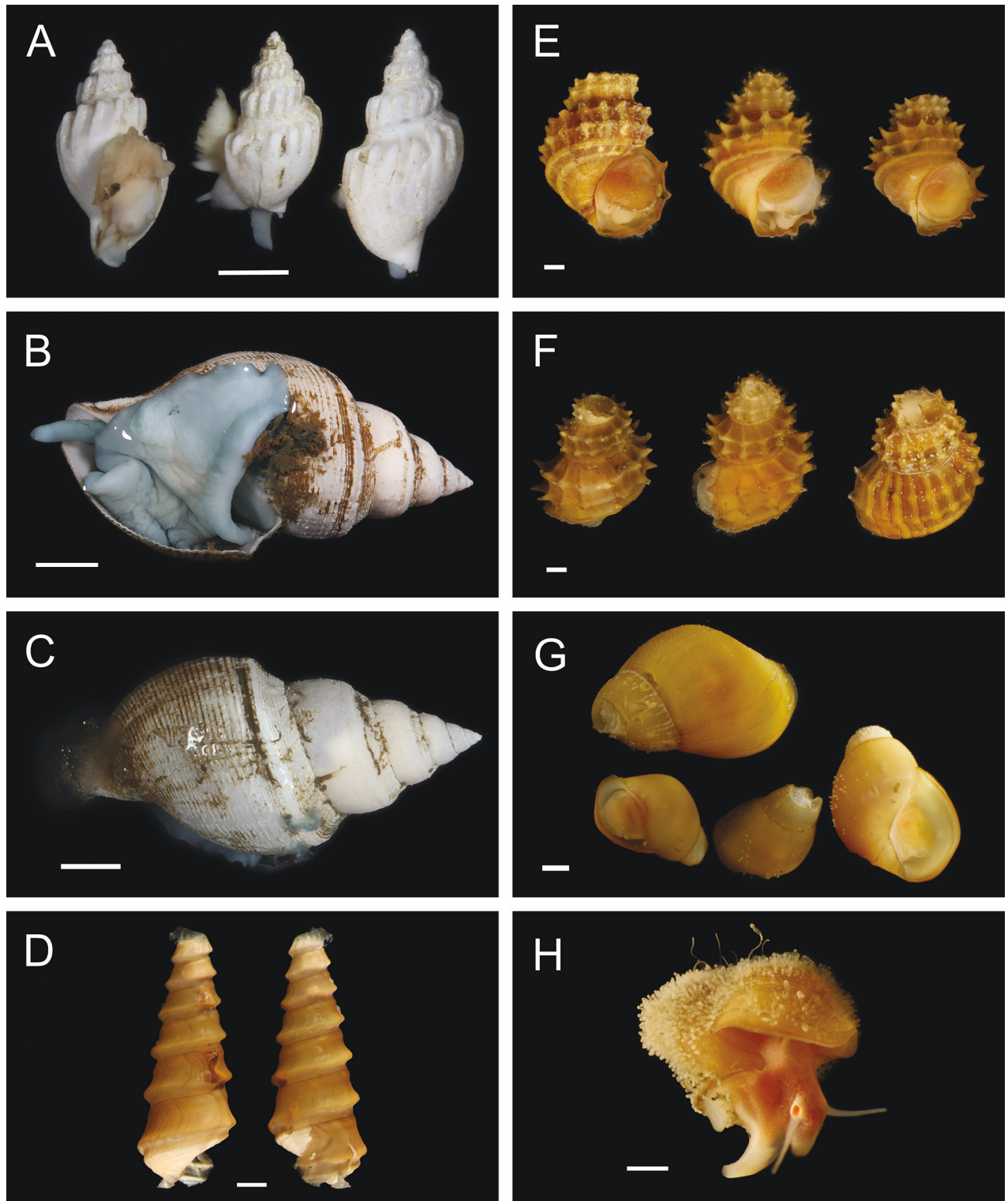


Figure 33. Mollusca: Gastropoda: Caenogastropoda, representative images. Live specimens are depicted unless otherwise specified **A** *Gymnobela* stet. (M17041, three specimens) **B** *Phymorhynchus* gen. inc. (M17038, apertural view) **C** *Phymorhynchus* gen. inc. (M17038, lateral view) **D** *Abysochrysos* stet. (M18994, same preserved specimen in lateral and apertural view) **E** *Provanna ios* (M16807, apertural views) **F** *Provanna ios* (M16807, apico-lateral views) **G** *Provanna laevis* (M16104) **H** *Provanna laevis* (M17030, with bacteria). Scale bars: 1 cm (**A–C**); 1 mm (**D–H**).

Mollusca | Gastropoda | Caenogastropoda | Abysochrysoidea |
Abysochrysidae

***Abysochrysos* stet.**

Fig. 33D

Material examined. AD4587: M18994.

Localities. Mound 12 (~ 9900–996 m).

Mollusca | Gastropoda | Caenogastropoda | Abysochrysoidea |
Provannidae

***Provanna ios* Warén & Bouchet, 1986**

Fig. 33E, F

Reference. Betters and Cordes 2024.

Localities. Jacó Scar (~ 1800–2000 m) and an unnamed locality (8.5958, -84.4370) at 1917 m depth, ca 12 km northwest of Parrita Seep.

Distribution. Eastern Pacific vents, 21°N to 17°S (type locality: 13°N on the East Pacific Rise) (Warén and Bouchet 1986), and the CRM seeps, 1757–2620 m depth (Sasaki et al. 2010; Betters and Cordes 2024).

***Provanna laevis* Warén & Ponder, 1991**

Fig. 33G, H

References. Betters and Cordes 2024.

Localities. Jacó Summit (~ 740–760 m), Mound 12 (~ 900–1050 m), The Thumb (~ 1071–1075 m).

Distribution. Originally described from the Guaymas Basin, Gulf of California, 2004 m (Warén and Ponder 1991), and reported at eastern Pacific vents and seeps from the Juan de Fuca Ridge to the CRM, 500–2004 m depth (Warén and Bouchet 2001; Betters and Cordes 2024). A recent synonymization of *Provanna glabra* Okutani, Tsuchida & Fujikura, 1992, extends the range of *P. laevis* to western Pacific seeps from Iheya Ridge to Sagami Bay (Betters and Cordes 2024).

Remarks. Some CRM specimens were associated with naturally occurring or experimentally deployed wood (Betters et al. 2023; Betters and Cordes 2024). Consistent with previous reports from the Oregon Margin (Warén and Bouchet 2001), *Pyropelta* cf. *corymba* was often attached to the shells. Some *P. laevis* shells were coated with elongated pustules (Fig. 33H) resembling the *Thiomargarita*-like sulfur-oxidizing bacteria reported on other gastropods from the CRM seeps (Bailey et al. 2011). Similar pustules have been observed on the shells of vent gastropods such as *Depressigyra* spp. (Warén and Bouchet 1989; Desbruyères et al. 2006) and *Lepetodrilus gordensis* S. B. Johnson, C. R. Young, W. J. Jones, Warén & Vrijenhoek, 2006 (Johnson et al. 2006).

***Provanna pacifica* (Dall, 1908)**

Fig. 34A, B

Reference. Betters and Cordes 2024.

Localities. Mound 11 (~ 1000–1100 m), Parrita Seep (~ 1300–1500 m).

Distribution. Associated with naturally occurring sunken wood and seeps, Oregon Margin to the Gulf of Panama (type locality), 1000–2750 m (Warén and Bouchet 1986; Betters and Cordes 2024).

Mollusca | Gastropoda | Heterobranchia | Euthyneura | Ringipleura |
Nudibranchia | Cladobranchia

Aeolidioidea stet.

Fig. 34C

Material examined. AD4975: M16811; AD4985: M16907.

Localities. Mound 12 (997–1002 m).

Fionoidea stet.

Fig. 34D

Material examined. AD4504: M11992; AD4906: M16094.

Localities. Mound 11 (~ 1004–1011 m), Mound 12 (~ 997–1002 m).

Remarks. M11992 was associated with naturally occurring sunken plant material.

Mollusca | Gastropoda | Heterobranchia | Euthyneura | Ringipleura |
Nudibranchia | Doridina

Goniodorididae sp. SIO_BIC_M16185

Fig. 34E, F

Material examined. AD4918: M16185 ([PQ449415](#)).

Localities. Quepos Slide (394 m).

Remarks. The closest COI BLASTN results on GenBank were within Goniodorididae, e.g., *Ceratodoris pilosa* (Bouchet & Ortea, 1983) from Japan ([MW357567.1](#), 88.94% identity), *Okenia mediterranea* (Ihering, 1886) from Italy ([MK645760.1](#), 88.78% identity), and *O. amoenula* (Bergh, 1907) from South Africa ([KF192606.1](#), 88.63% identity).

Mollusca | Gastropoda | Heterobranchia | Euthyneura | Tectipleura |
Pyramidellidae

***Eulimella lomana* (Dall, 1908)**

Fig. 34G

Reference. Electronic supplemental table S6 of Levin et al. (2012) (occurrences only, Jacó Scar).

Material examined. AD4590: M18971.

Localities. Jacó Scar (~ 1800 m).

Distribution. Known from seeps, vents (likely sedimented vents), and whale falls, 1168–2008 m, from southern California (type locality) and the Guaymas Basin, Gulf of California (Warén and Bouchet 1993; Sasaki et al. 2010; Portail et al. 2015).

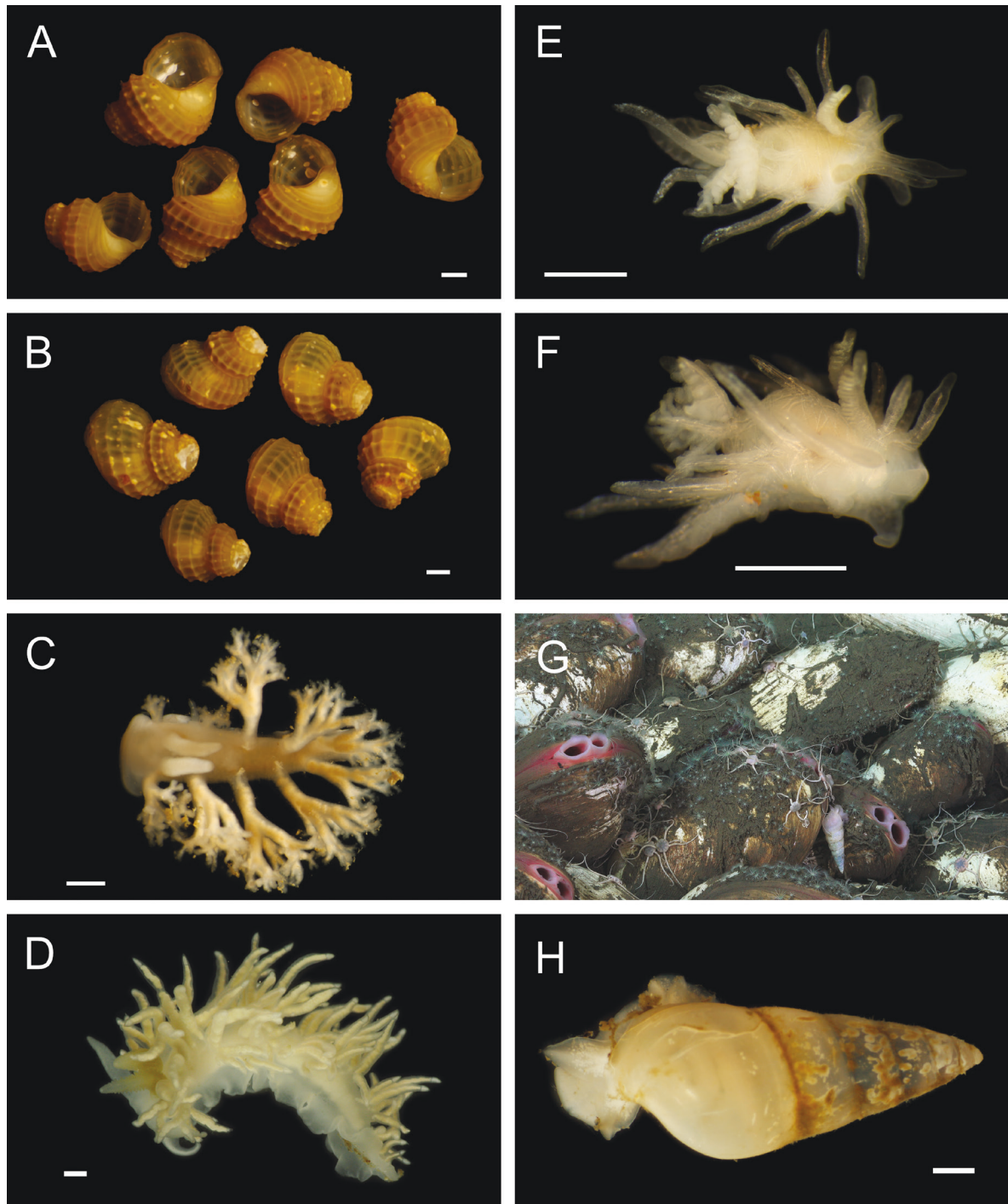


Figure 34. Mollusca: Gastropoda: Caenogastropoda and Heterobranchia, representative live images **A** *Provanna pacifica* (M16955, apertural views) **B** *Provanna pacifica* (M16955, lateral views) **C** *Aeolidioidea* stet. (M16811) **D** *Fionoidea* stet. (M11992) **E** *Goniodorididae* sp. SIO_BIC_M16185 (M16185, dorsal view) **F** *Goniodorididae* sp. SIO_BIC_M16185 (M16185, lateral view) **G** *Eulimella lomana* (*in situ*), Dive S0214 at Jacó Scar, 1781 m. Credit: ROV SuBastian/Schmidt Ocean Institute **H** *Pyramidellidae* stet. (M16901). Scale bars: 1 mm.

New records. The CRM specimen represents a new southern record for this species.

Remarks. Pyramidellids are thought to be exclusively parasitic, typically on mollusks or polychaetes, but a definitive host has not been identified for *E. lomana* (Warén and Bouchet 1993). Specimen M18971 could not be linked to

a potential host. Another specimen at Jacó Scar (not collected) was observed on vesicomylid clams (Fig. 34G).

Pyramidellidae stet.

Fig. 34H

Material examined. AD4508: M12029; AD4912: M16125; AD4987: M16901; AD4989: M16942.

Localities. Mound 12 (1010 m), Parrita Seep (~ 1401–1419 m), Jacó Scar (1842 m).

Remarks. At least one additional morphospecies is present and distinct from *E. lomana*.

Mollusca | Gastropoda | Heterobranchia | Mesoneura | Tjaernoioidea | Aplustridae

Parvaplustrum stet.

Fig. 35A, B

Material examined. S0219: M17071.

Localities. Rio Bongo Scar (609 m).

Remarks. Collected from a microbial mat. DNA sequences could not be obtained.

Mollusca | Gastropoda | Heterobranchia | “Lower Heterobranchia” | Architectonicoidea | Architectonicidae

Architectonicidae fam. inc.

Fig. 35C

Material examined. AD4505: M19335.

Localities. Mound 11 (1025 m).

Remarks. Identification is uncertain due to the corroded condition of the shell.

Mollusca | Gastropoda | Heterobranchia | “Lower Heterobranchia” | Orbitestelloidea | Orbitestellidae

Lurifax gen. inc.

Fig. 35D

Material examined. AD4504: M19337; AD4505: M19336.

Localities. Mound 11 (1009–1025 m).

Remarks. Identification is uncertain.

Orbitestella gen. inc.

Fig. 35E

Material examined. AD4589: M19136.

Localities. Mound 12 (997 m).

Remarks. Identification is uncertain due to the corroded condition of the shell.

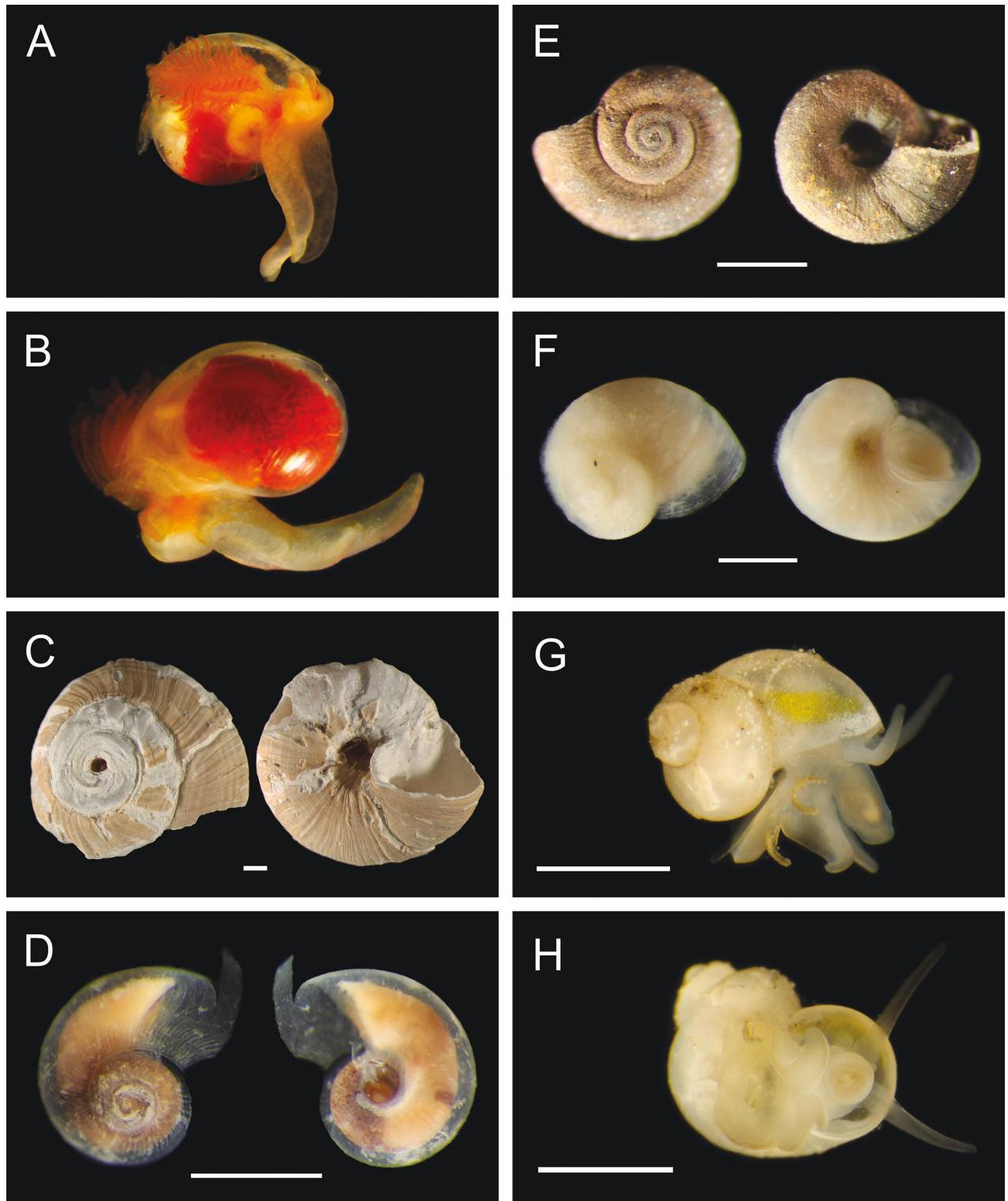


Figure 35. Mollusca: Gastropoda: Heterobranchia, representative images. Live specimens are depicted unless otherwise specified **A** *Parvaplustrum* stet. (M17071, apertural view; scale not recorded for specimens destroyed in DNA extraction; estimated shell length 1-2 mm) **B** *Parvaplustrum* stet. (M17071, lateral view) **C** Architectonicidae fam. inc. (M19335, same preserved specimen in apical and umbilical view) **D** *Lurifax* gen. inc. (M19337, same preserved specimen in apical and umbilical view) **E** *Orbitestella* gen. inc. (M19136, same preserved specimen in apical and umbilical view) **F** *Hyalogyra* stet. (M18944, same preserved specimen in apical and umbilical view) **G** *Hyalogyrina* sp. SIO_BIC_M16774 (M16774, lateral view) **H** *Hyalogyrina* sp. SIO_BIC_M16774 (M16774, ventral view). Scale bars: 1 mm.

Mollusca | Gastropoda | Heterobranchia | “Lower Heterobranchia” |
Valvatoidea | Hyalogyrinidae

***Hyalogyra* stet.**

Fig. 35F

Material examined. AD4586: M18799; AD4587: M19010, M19087; AD4588: M19026, M19043, M19074, M19088, M19135; AD4589: M18837, M19137; AD4590: M18907, M18919, M18920, M18942, M18944, M19145, M19338.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. Two morphospecies were distinguished as “tall” and “planispiral”.

***Hyalogyrina* stet.**

Fig. 35G, H

Reference. Electronic supplemental table S6 of Levin et al. (2012) (occurrences only, Jacó Scar).

Material examined. AD4510: M12043; AD4587: M13003, M13004, M18991, M18992, M19000, M19001, M19007, M19144, M19147; AD4590: M18323, M18981, M19330; AD4914: M16151 ([PQ449414](#)); AD4974: M16774; S0213: M17029 ([PQ449419](#)).

Localities. Jacó Summit (741–742 m), Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. Most specimens were collected from microbial mat habitats. At least two morphospecies were distinguished (Levin et al. 2012). One morphospecies, represented by M17029, has a pale shell with a broad yellow blaze on the outer whorl. The CRM specimens warrant comparison to *Hyalogyrina grasslei* Warén & Bouchet, 1993, which is known from hydrothermal vents and seeps at ~ 2000 m in the Guaymas Basin, Gulf of California (Warén and Bouchet 1993; Sasaki et al. 2010).

Mollusca | Gastropoda | Vetigastropoda | Lepetellida | Lepetelloidea |
Caymanabyssiidae

***Colotrachelus* stet.**

Fig. 36A

Material examined. AD4587: M19093, M19101, M19120, M19129.

Localities. Mound 12 (~ 1000 m).

Mollusca | Gastropoda | Vetigastropoda | Lepetellida | Lepetelloidea |
Pyropeltidae

***Pyropelta* cf. *corymba* McLean & Haszprunar, 1987**

Fig. 36B

Reference. Reported in Electronic supplemental table S6 of Levin et al. (2012) as *Pyropelta corymba* (occurrences only, Jacó Scar).

Material examined. AD4501: M11968; AD4586: M18801; AD4587: M19096, M19343; AD4588: M19033, M19049, M19059, M19073, M19160, M19347; AD4589: M18322, M18830, M18841, M18851, M18863, M19078; AD4590: M18908; AD4910: M16105, M16107; AD4917: M16157; AD4978: M16874.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. *Pyropelta corymba* was originally described from hydrothermal vents in the Guaymas Basin, Gulf of California, 2022 m (McLean and Haszprunar 1987), and subsequently reported from seeps on the Oregon Margin, 524 m (Warén and Bouchet 2001), as well as from whale falls off southern California, 940–1240 m (McLean 1992). If confirmed as *P. corymba* by genetic comparison to material from the type locality, the CRM seep specimens would represent new southern records for this species. Consistent with previous reports (Warén and Bouchet 2001; Desbruyères et al. 2006), many CRM specimens were associated with *Provanna laevis*.

***Pyropelta* cf. *musaica* McLean & Haszprunar, 1987**

Fig. 36C

Material examined. AD4586: M17850, M18781, M18811, M19348; AD4587: M19008, M19090, M19109, M19113, M19122; AD4588: M19022, M19023, M19032, M19044, M19054, M19066, M19085, M19162, M19346; AD4589: M18831, M18843, M18850, M19077; AD4590: M18940, M18988; AD4591: M18324, M18883, M18900.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. *Pyropelta musaica* was originally described from hydrothermal vents at the Axial Seamount, Juan de Fuca Ridge, 1575 m (McLean and Haszprunar 1987). It has been reported from the Jalisco Block seeps, Mexico, at 3000–3775 m (Warén and Bouchet 2001; Desbruyères et al. 2006), and from whale falls across California at 940–1400 m (McLean 1992). If confirmed as *P. musaica* by genetic comparison to material from the type locality, the CRM seep occurrences would represent new southern records for this species.

***Pyropelta* cf. *wakefieldi* McLean, 1992**

Fig. 36D, E

Material examined. AD4501: M11966 ([PQ450396](#)); AD4586: M17849, M18797, M18803, M18808, M18810; AD4587: M19011, M19016, M19112; AD4588: M19021, M19030, M19048, M19072, M19075, M19134, M19148, M19150, M19340, M19341, M19342, M19344; AD4589: M18829, M18835, M18849, M18857, M18862, M19076; AD4590: M18956, M18959.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. *Pyropelta wakefieldi* was originally described from a whale fall off Point Sur, California, 940 m (McLean 1992). Genetic confirmation of the CRM seep occurrences would represent new southern records and new seep records for this species. The closest COI BLASTN result on GenBank was *Pyropelta* sp. SWA-2009 from the Gulf of Mexico ([FJ977753.1](#), 90.99% identity).

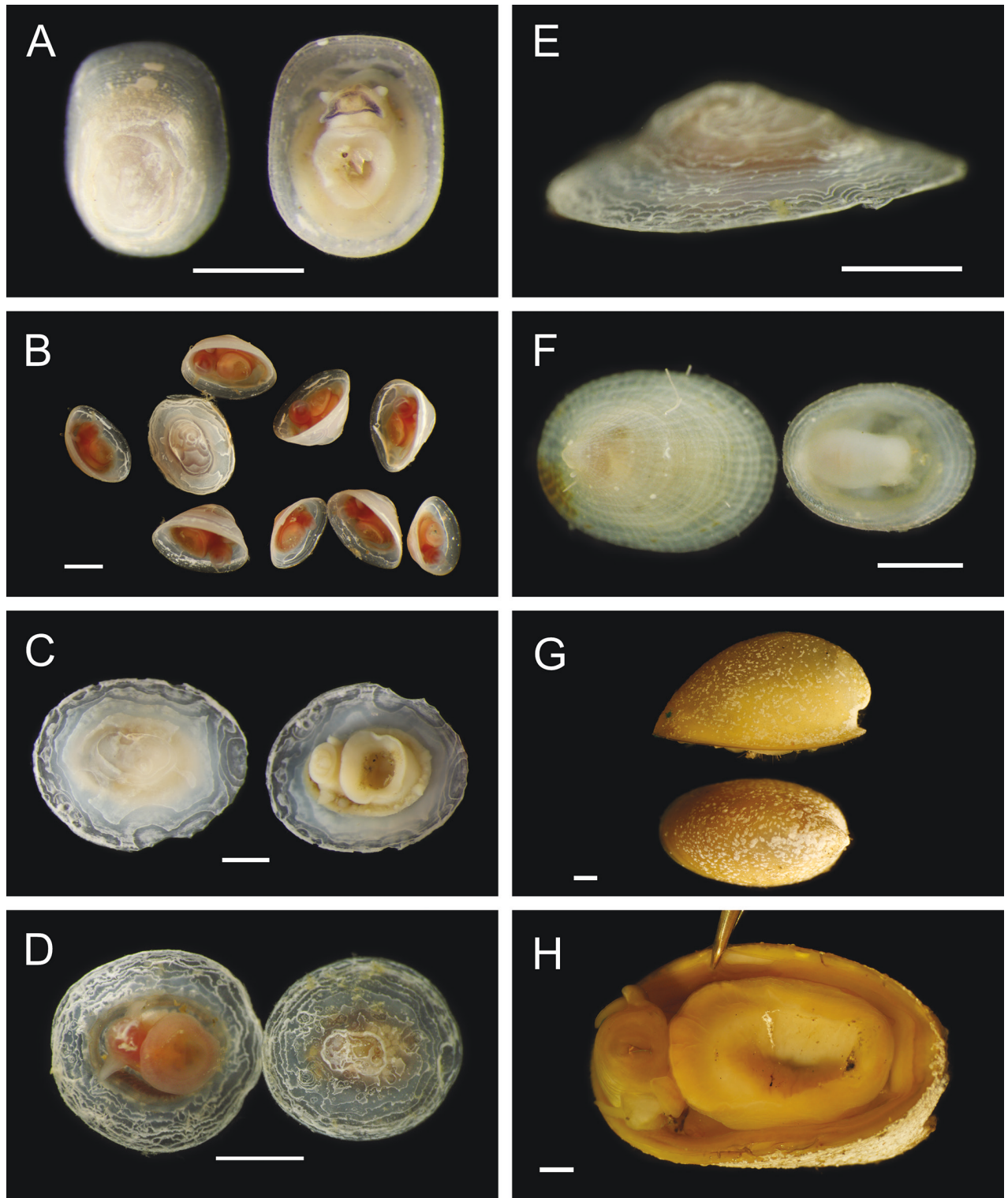


Figure 36. Mollusca: Gastropoda: Vetigastropoda, representative images. Live specimens are depicted unless otherwise specified **A** *Colotrachelus stet.* (M19101, separate preserved specimens in dorsal and ventral view) **B** *Pyropelta cf. corymba* (M16105) **C** *Pyropelta cf. musaica* (M19044, same preserved specimen in dorsal and ventral view) **D** *Pyropelta cf. wakefieldi* (M11966, separate specimens in dorsal and ventral view) **E** *Pyropelta cf. wakefieldi* (M11966, lateral view) **F** *Lepetodrilus aff. shannonae* (M11973, separate specimens in dorsal and ventral view) **G** *Lepetodrilus guaymasensis* (M16142, separate specimens in lateral and dorsal view) **H** *Lepetodrilus guaymasensis* (M16142, ventral view). Scale bars: 1 mm.

Mollusca | Gastropoda | Vetigastropoda | Lepetellida | Lepetodrilioidea |
Lepetodrilidae

***Lepetodrilus* aff. *shannonae* Warén & Bouchet, 2009**

Fig. 36F

Reference. Electronic supplemental table S6 of Levin et al. (2012) (occurrences only, Jacó Scar).

Material examined. AD4501: M11973; AD4511: M12057; AD4590: M18904, M18943, M18985, M19152; AD4591: M18875, M18879, M18891, M18898.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. The CRM specimens appear morphologically similar to *L. shannonae*, known only from hydrocarbon seeps off the Congo River, West Africa, 2300–3150 m (Warén and Bouchet 2009).

***Lepetodrilus* *guaymasensis* McLean, 1988**

Fig. 36G, H

References: Johnson et al. 2008; Matabos and Jollivet 2019; Electronic supplemental table S6 of Levin et al. (2012) (occurrences only, Jacó Scar).

Additional material examined. AD4508: M12020 ([PQ450397](#)); AD4586: M18778, M18809; AD4587: M19107, M19111, M19118; AD4588: M19018, M19028, M19058, M19069; AD4589: M18817, M18828, M18836, M18845, M18856, M18860; AD4590: M18911, M18914, M18922, M18927, M18938, M18947, M18950, M18967, M18970, M18979, M18983, M19349 to M19382; AD4591: M18872, M18878, M18890, M18897; AD4912: M16142; AD4917: M16159.

Localities. Mound 12 (~ 1000 m; this study), Parrita Seep (~ 1400 m; this study), Jacó Scar (~ 1800 m), “Mudpie” seep ~ 10 km west of Parrita Scar (8.983, -84.717; 1917 m) (Johnson et al. 2008).

Distribution. Originally described from sedimented hydrothermal vents and seeps in the Guaymas Basin, Gulf of California, 2000–2019 m, often in association with *Riftia pachyptila* Jones, 1981 (McLean 1988, 1993; Sasaki et al. 2010). Specimens from the CRM “Mudpie” seep were confirmed as *L. guaymasensis* based on COI comparison to specimens from the type locality (Matabos and Jollivet 2019).

Remarks. The COI sequence of M12020 was 100.00% identical to that of *L. guaymasensis* from the CRM Mudpie site ([EU306419.1](#), voucher SMNH 82443, Swedish Museum of Natural History).

Lepetodrilus* *stet.

Fig. 37A

Material examined. AD4501: M11965.

Localities. Mound 12 (~ 984–997 m).

Remarks. Additional *Lepetodrilus* morphospecies may occur at the CRM. For example, an undescribed *Lepetodrilus* “sp. CR” has been reported from a CRM seep site at 1900 m, but it is represented by a single specimen without available DNA sequences (Johnson et al. 2008). Specimen M11965 and others resemble *L. elevatus* J. H. McLean, 1988, which was originally described as two subspecies from the East Pacific Rise and the Galápagos Rift (McLean 1988).

The *L. elevatus* species complex comprises four genetically distinct lineages (Johnson et al. 2008; Matabos and Jollivet 2019). Additional specimens of Lepetodrilidae from the CRM are under genetic and morphological investigation (Betters et al. (2024), in press at the time of this work's acceptance).

Mollusca | Gastropoda | Vetigastropoda | Lepetellida | Scissurelloidea | Anatomidae

Anatoma stet.

Fig. 37B

Reference. Electronic supplemental table S6 of Levin et al. (2012) (occurrences only, Jacó Scar).

Material examined. AD4591: M18899; AD4918: M16187.

Localities. Quepos Slide (~ 333–408 m), Jacó Scar (1753 m).

Mollusca | Gastropoda | Vetigastropoda | Lepetellida | Scissurelloidea | Larocheidae

Bathyxylophila stet.

Fig. 37C, D

Material examined. AD4508: M12028; AD4509: M12038, M12039, M18869; AD4587: M18317, M19092, M19099, M19128; AD4923: M16198, M16199; AD4976: M16797, M16798, M16801; AD4988: M16954, M16965.

Localities. Mound 12 (~ 1000 m), Mound 11 (1010 m), Parrita Seep (~ 1000–1400 m), Jacó Scar (~ 1800 m).

Remarks. Associated with naturally occurring or experimentally deployed wood falls. Likely at least two morphospecies are represented. Some of the specimens of M16199 (Fig. 37D) require further verification.

Mollusca | Gastropoda | Vetigastropoda | Seguenziida | Seguenzioidea | Cataegidae

Kanoia myronfeinbergi Warén & Rouse, 2016

Fig. 37E

References. Warén and Rouse 2016**. Prior to description, this morphospecies was reported at Jacó Scar as “*Cataegis* sp.” in Electronic supplemental table S6 of Levin et al. (2012).

New sequences. We provide COI sequences for the following specimens cited in Warén and Rouse (2016), in some cases under catalog numbers from the Swedish Museum of Natural History, SMNH: AD4501: M11976 ([PQ449395](#)); AD4587: M14422 ([PQ449410](#); ex SMNH 108692), M14424 ([PQ449411](#); ex SMNH 109248); AD4588: M18331 ([PQ449427](#); ex SMNH 108742); AD4589: M14418 ([PQ449405](#); tissue from voucher SMNH 108441); AD4590: M14421 ([PQ449409](#); tissue from voucher SMNH 108604); AD4591: M14419 ([PQ449406](#); ex SMNH 108504), M14420A ([PQ449407](#); ex SMNH 108525), M14420B ([PQ449408](#); ex SMNH 108525); Guaymas Basin seeps: M13149

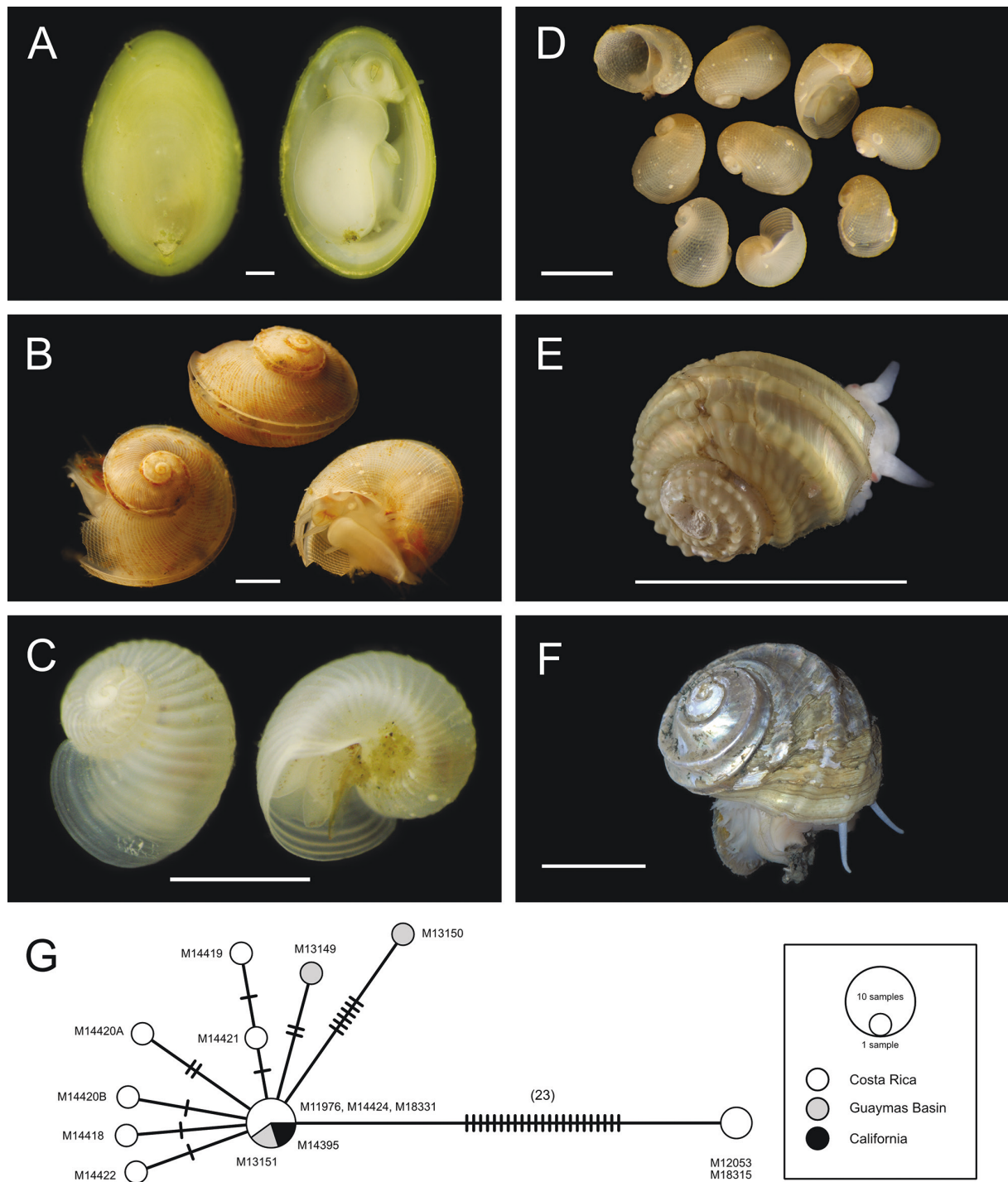


Figure 37. Mollusca: Gastropoda: Vetigastropoda, representative live images **A** *Lepetodrilus stet* (M11965, same specimen in dorsal and ventral view) **B** *Anatoma stet*. (M16187, same specimen in apical, lateral, and umbilical view) **C** *Bathyxylophila stet*. (M12038, separate specimens in apical and umbilical view) **D** *Bathyxylophila stet*. (M16199) **E** *Kanoia myronfeinbergi* (M16771) **F** *Kanoia cf. myronfeinbergi* (M12053) **G** Haplotype network of *Kanoia* COI sequences. Scale bars: 1 mm (A–D); 1 cm (E, F).

([PQ432664](#)), M13150 ([PQ432665](#)), M13151 ([PQ432666](#)); Del Mar seeps: M14395 ([PQ432667](#)).

Localities. Mound 12 (~ 1000 m; type locality), Jacó Scar (~ 1800 m); additional seep sites off Costa Rica and Nicaragua (1002–1917 m) (Warén and Rouse 2016).

Distribution. Also known from seeps in the Guaymas Basin at ~ 1570 m and seeps off Del Mar, California, ~ 1020 m (Warén and Rouse 2016).

***Kanoia* cf. *myronfeinbergi* Warén & Rouse, 2016**

Fig. 37F

Reference. Warén and Rouse 2016.

New sequences. AD4510: M12053 ([PQ449403](#)); AD4587: M18315 ([PQ449426](#); tissue from voucher SMNH 108680). In Warén and Rouse (2016), specimen M12053 was morphologically identified as *K. myronfeinbergi* whereas M18315 was not identified to species due to corrosion of the shell.

Localities. Jacó Summit (~ 741–744 m), Mound 12 (~ 990–996 m).

Remarks. The description of *K. myronfeinbergi* notes the possibility of a second, cryptic species “with less distinct sculpture, more similar to *Kanoia mero-glypta* from the Caribbean,” although detailed assessment of shell sculpture is difficult due to the corrosion on many specimens (Warén and Rouse 2016).

To investigate the genetic basis for this variation, we constructed a COI haplotype network using a subset of the specimens examined in the original description (Fig. 37G). Two COI sequences showed a minimum uncorrected distance of 4.3% (over 539 bp) from the others. Although there is no universally applicable threshold for species delimitation, numerous deep-sea gastropod species are separated by a minimum of 4–7% COI divergence (Johnson et al. 2008; Chen et al. 2019a). We conservatively designate the two high-divergence specimens as *K. cf. myronfeinbergi*, and we note the co-occurrence of this haplotype with others at Mound 12. The other *K. myronfeinbergi* sequences showed minimal variation across localities from Costa Rica to California (maximum uncorrected distance 1.11%). Further clarification on *Kanoia* species delimitation will require genetic and morphological analysis of more individuals from various localities and depths, especially those < 1000 m to explore potential depth segregation.

Mollusca | Gastropoda | Vetigastropoda | Seguenziida

***Xyloskenea* stet.**

Fig. 38A

Material examined. AD4509: M18870, M19159; AD4587: M19013, M19334; AD4923: M16197.

Localities. Mound 12 (~ 1000 m), Parrita Seep (~ 1100 m), Jacó Scar (~ 1800 m).

Remarks. M16197 and M18870 were associated with naturally occurring wood falls.

Mollusca | Gastropoda | Vetigastropoda | Trochida | Trochoidea | Colloniidae

***Escondidacantanea panamensis* (Dall, 1908)**

Fig. 38B, C

Material examined. AD4587: M17848, M19103.

Localities. Mound 12 (995–996 m).

Distribution. Originally described from the Gulf of Panama, 1015 m (Dall 1908), and recorded south to the Concepción seeps off central Chile, 740–870 m (Sasaki et al. 2010).

Remarks. M17848 and M19103 were associated with naturally occurring wood falls.

Mollusca | Gastropoda | Vetigastropoda | Trochida | Trochoidea | Skeneidae

***Dillwynella* stet.**

Fig. 38D

Reference. Reported in Electronic supplemental table S6 of Levin et al. (2012) as “*Dillwynella panamensis*” (occurrences only, Jacó Scar).

Material examined. AD4586: M18805; AD4587: M18320, M19095, M19098, M19117, M19124, M19158; AD4588: M17884, M19052; AD4589: M18832; AD4590: M18957.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. Many specimens were associated with wood falls, either experimentally deployed (M17884, M18805, M19052) or naturally occurring (M18320, M19095, M19098, M19117, M19124, M19158). These specimens warrant comparison to *Ganesa panamensis* Dall, 1902, which is known only from the Gulf of Panama, 1865 m, and has been regarded as possibly belonging to the genus *Dillwynella* (Dall 1902; Kunze 2011).

***Fucaria* stet.**

Fig. 38E

Reference. Electronic supplemental table S6 of Levin et al. (2012) (occurrences only, Jacó Scar).

Material examined. AD4513: M12068; AD4591: M18880, M18887; AD4590: M18906, M18923, M18941, M18946, M18951, M18986, M19165; AD4591: M18896.

Localities. Jacó Scar (~ 1800 m).

Mollusca | Gastropoda | Neomphaliones

We follow the subdivisions of Neomphaliones based on the mitogenome phylogeny in Zhong et al. (2022).

Mollusca | Gastropoda | Neomphaliones | Cocculinida | Bathysciadiidae

***Bathysciadium* stet.**

Fig. 38F

Material examined. AD4587: M19002.

Localities. Mound 12 (~ 990–996 m).

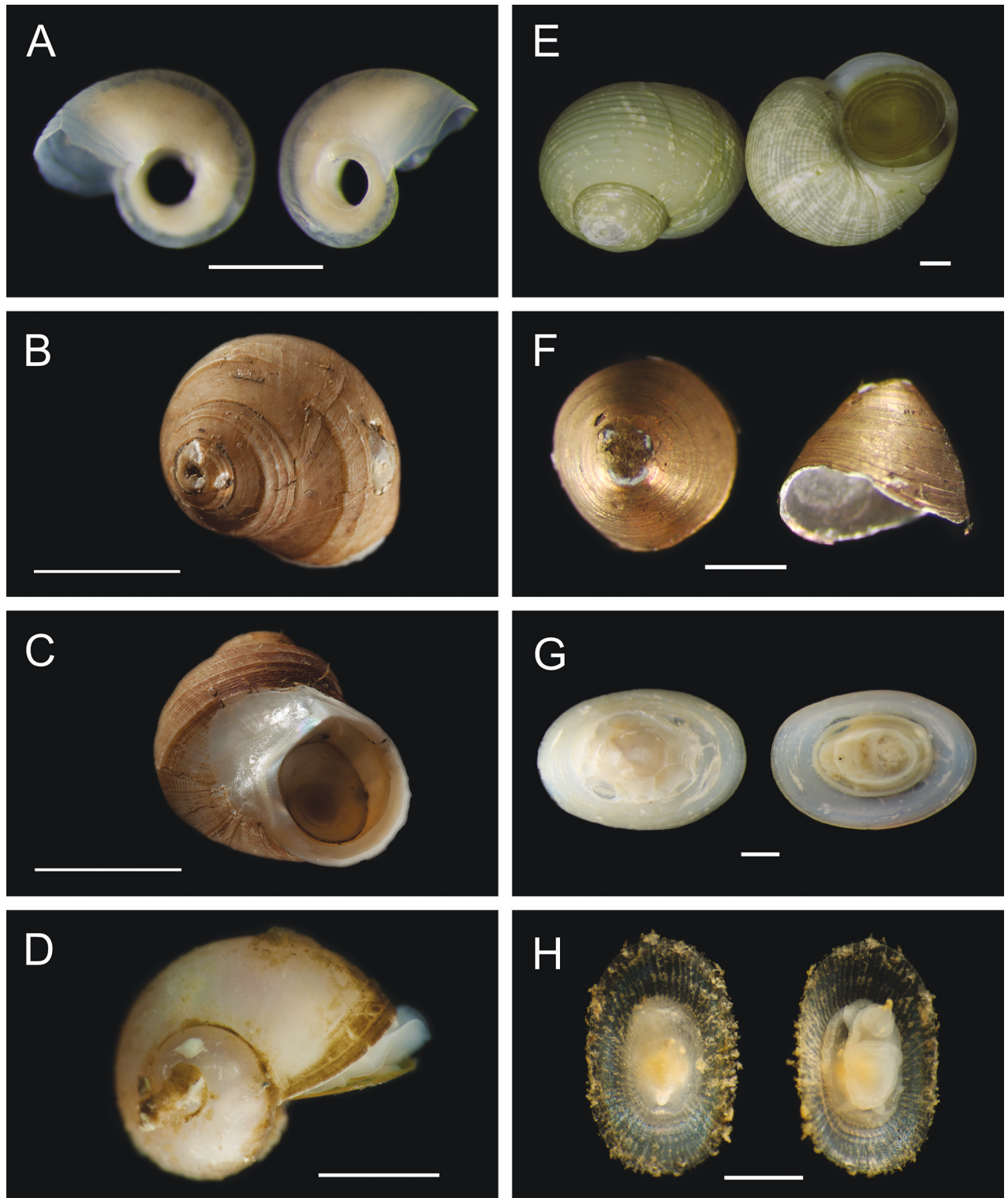


Figure 38. Mollusca: Gastropoda: Vetigastropoda and Neomphaliones, representative images. Live specimens are depicted unless otherwise specified **A** *Xyloskenea* stet. (M19013, two preserved specimens) **B** *Escondidacantainea panamensis* (M17848, apical view) **C** *Escondidacantainea panamensis* (M17848, umbilical view) **D** *Dillwynella* stet (M17884) **E** *Fucaria* stet. (M12068, two specimens) **F** *Bathysciadium* stet. (M19002, same preserved specimen in dorsal and lateral view) **G** *Cocculina* stet. (M19166, same preserved specimen in dorsal and ventral view) **H** Cocculinidae fam. inc. (M16727, same specimen in dorsal and ventral view). Scale bars: 1 mm (**A**, **D–H**); 1 cm (**B**, **C**).

Mollusca | Gastropoda | Neomphaliones | Cocculinida | Cocculinidae

***Cocculina* stet.**

Fig. 38G

Material examined. AD4503: M19166; AD4588: M18321, M19089.

Localities. Mound 12 (~ 1000 m).

Cocculinidae fam. inc.

Figs 38H, 39A, B

Material examined. AD4972: M16726, M16727, M16728; AD4974: M16789, M16790; AD4976: M16816, M16817.

Localities. Mound 12 (~ 1000 m), Jacó Scar (~ 1800 m).

Remarks. Associated with experimentally deployed wood. Under morphological and genetic investigation (Betters et al. (2024), in press at the time of this work's acceptance).

Mollusca | Gastropoda | Neomphaliones | Neomphalida |
Melanodrymiidae

***Leptogyropsis* gen. inc.**

Fig. 39C

Material examined. AD4587: M19102, M19146.

Localities. Mound 12 (996 m).

Remarks. Associated with a naturally occurring wood fall.

Mollusca | Gastropoda | Neomphaliones | Neomphalida |
Neomphalidae

***Helicrenion* stet.**

Fig. 39D

Material examined. AD4508: M12027; AD4586: M18782; AD4587: M18993, M18999, M19004; AD4589: M18818; AD4590: M18909; AD4988: M16962.

Localities. Mound 11 (1009 m), Mound 12 (~ 1000 m), Parrita Seep (~ 1400 m), Jacó Scar (~ 1800 m).

Remarks. Possibly an undescribed species. Several specimens (M16962, M18993, M18999, and M19004) were collected from microbial mat habitats.

Neomphalidae sp. SIO_BIC_M14645

Fig. 39E, F

Material examined. AD4910: M14645; AD4917: M14646 ([PQ449412](#)).

Localities. Mound 12 (~ 1000 m).

Remarks. An undescribed genus and species.

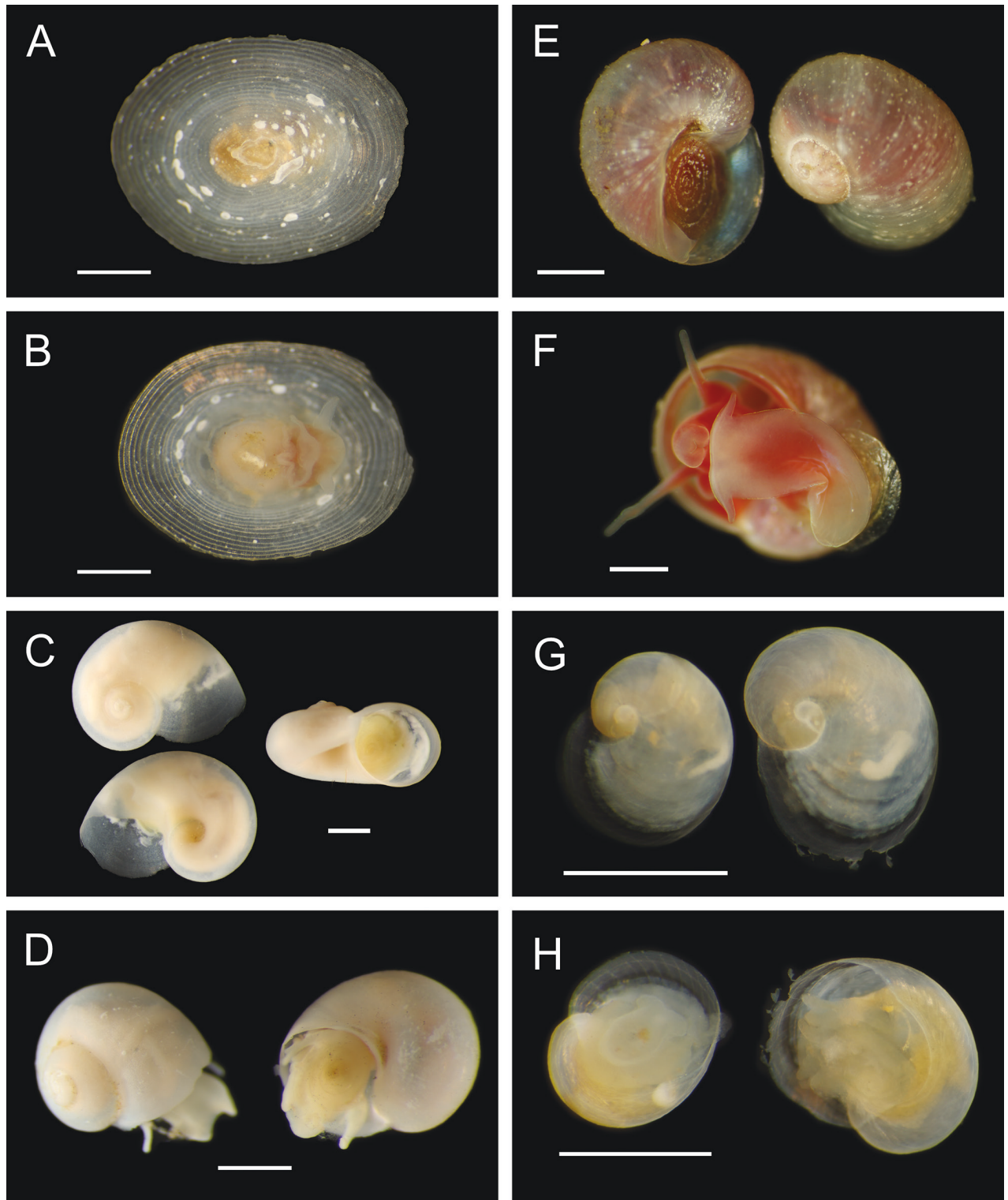


Figure 39. Mollusca: Gastropoda: Neomphaliones, representative images. Live specimens are depicted unless otherwise specified **A** Cocculinidae fam. inc. (M16816, dorsal view) **B** Cocculinidae fam. inc. (M16816, ventral view) **C** *Leptogyropsis* gen. inc. (M19102, same preserved specimen in apical, umbilical, and apertural view) **D** *Helicrenion* stet. (M18909, two preserved specimens) **E** Neomphalidae sp. SIO_BIC_M14645 (M14645, two specimens) **F** Neomphalidae sp. SIO_BIC_M14645 (M14645, ventral view) **G** Peltospiridae stet. (M16802, dorsal view) **H** Peltospiridae stet. (M16802, ventral view). Scale bars: 1 mm.

Mollusca | Gastropoda | Neomphaliones | Neomphalida | Peltospiridae

Peltospiridae *stet.*

Fig. 39G, H

Material examined. AD4976: M16802.

Localities. Jacó Scar (1887 m).

Remarks. Associated with experimentally deployed wood.

Mollusca | Scaphopoda

Gadilida *stet.*

Fig. 40A

Material examined. AD4510: M12049.

Localities. Jacó Summit (744 m).

Siphonodentalium *gen. inc.*

Fig. 40B

Material examined. AD4510: M12048; AD4511: M12056.

Localities. Jacó Summit (742 m), Mound 12 (~ 988–997 m).

Mollusca | Cephalopoda

We thank Michael Vecchione (National Marine Fisheries Service National Systematics Laboratory, U.S. National Museum of Natural History) for assistance with these identifications.

Mollusca | Cephalopoda | Octopoda

Octopodoidea *sp.* **SIO_BIC_ M17037**

Fig. 40C, D

Material examined. S0216: M17037.

Localities. Quepos Slide (317 m).

Remarks. Likely an undescribed species, possibly an undescribed genus. The animal was observed releasing ink (potentially a diagnostic character).

Mollusca | Cephalopoda | Oegopsida

Planctoteuthis danae (Joubin, 1931)

Fig. 40E

Material examined. S0215: M17039.

Localities. Mound 12 (1016 m depth, 2–3 m above the seafloor).

Distribution. Originally described from the Gulf of Panama and considered cosmopolitan in tropical and temperate waters worldwide (Jereb and Roper 2010).

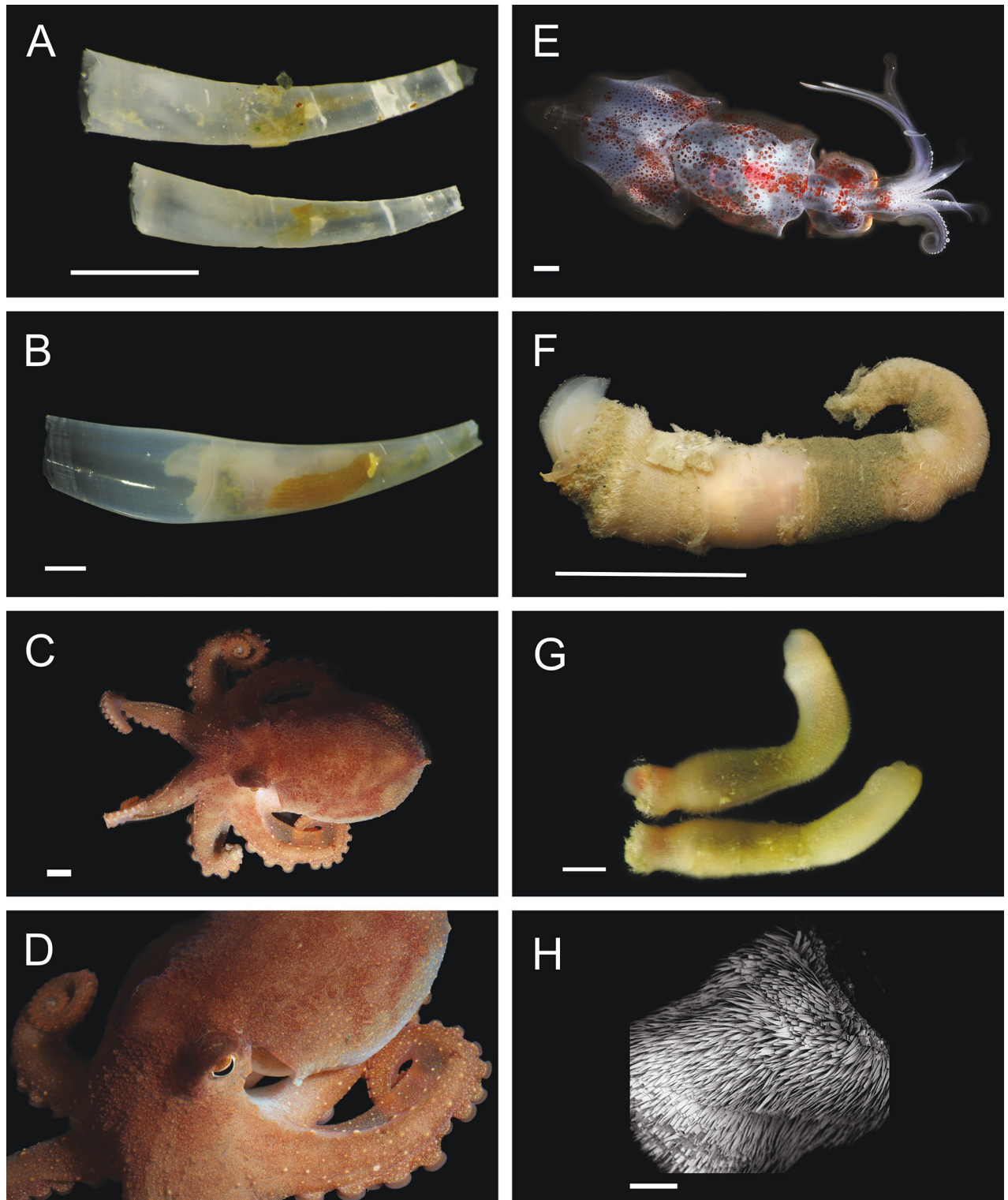


Figure 40. Mollusca: Scaphopoda, Cephalopoda, and Aplacophora: Caudofoveata, representative live and SEM images **A** *Siphonodentalium* gen. inc. (M12048) **B** *Gadilida* stet. (M12049) **C** *Octopodoidea* stet. (M17037, dorsal view) **D** *Octopodoidea* stet. (M17037, detail) **E** *Planctoteuthis danae* (M17039) **F** *Chaetoderma* stet. (M16155, incomplete specimen, cuticle missing from part of the body) **G** *Chaetodermatidae* sp. SIO_BIC_M12018 (M12018) **H** *Chaetodermatidae* sp. SIO_BIC_M12018 (M12019, SEM). Scale bars: 1 mm (**A**, **B**, **G**); 1 cm (**C**, **E**, **F**); 0.3 mm (**H**).

Mollusca | Caudofoveata

We use the clade-based high-level taxonomic names in Kocot et al. (2019).

***Chaetoderma* stet.**

Fig. 40F

Material examined. AD4917: M16155; AD4977: M16809.

Localities. Jacó Scar (1783–1791 m).

Chaetodermatidae sp. SIO_BIC_M12018

Fig. 40G, H

Material examined. AD4508: M12018 ([PQ449402](#)), M12019; AD4972: BI1338 ([PQ449340](#)).

Localities. Parrita Seep (~ 1401–1419 m), Jacó Scar (1746 m).

Remarks. The closest COI BLASTN result on GenBank was the holotype of *Chaetoderma felderi* Ivanov & Scheltema, 2007 ([AM922259.1](#); 93.33% identity to M12018, 93.18% identity to BI1338). Based on COI, these specimens belong within Chaetodermatidae and may belong to *Chaetoderma* or *Falcidens*, but these genera are not reciprocally monophyletic (Mikkelsen et al. 2019).

Chaetodermatidae sp. SIO_BIC_M16812

Material examined. AD4975: M16812 ([PQ449416](#); no image available).

Localities. Mound 12 (1000 m).

Remarks. The closest COI BLASTN result on GenBank was the holotype of *Chaetoderma felderi* ([AM922259.1](#); 90.21% identity). As above, this specimen may belong to *Chaetoderma* or *Falcidens*.

Chaetodermatidae sp. SIO_BIC_M16891

Fig. 41A

Material examined. AD4979: M16891 ([PQ435556](#)).

Localities. Quepos Slide (397 m).

Remarks. The closest COI BLASTN result on GenBank was an undescribed species of *Falcidens* ([MG855756.1](#); 93.93% identity). As above, this specimen may belong to *Chaetoderma* or *Falcidens*.

Mollusca | Solenogastres

We use the clade-based high-level taxonomic names in Kocot et al. (2019). Groups that have been historically assigned to the non-monophyletic “Cavibelonia” are listed last.

Mollusca | Solenogastres | Neomeniamorpha | Neomeniidae

***Neomenia* gen. inc.**

Fig. 41B

Material examined. S0219: M18409.

Localities. Rio Bongo Scar (606 m).

Mollusca | Solenogastres | Pholidoskepia | Gymnomeniidae

***Gymnomeniidae* stet.**

Fig. 41C, D

Material examined. AD4990: M16924 ([PQ435558](#); 16S: [PQ304664](#)).

Localities. Parrita Seep (1401 m).

***Wirenia* sp. SIO_BIC_M17072**

Fig. 41E, F

Material examined. S0219: M17072 ([PQ435553](#); 16S: [PQ304665](#)).

Localities. Rio Bongo Scar (606 m).

Remarks. This undescribed species has keeled, leaf-like sclerites typical of *Wirenia*, but it is easily distinguished from all described species by the extremely small size (< 200 µm) of the sclerites.

Mollusca | Solenogastres | Pholidoskepia

***Pholidoskepia* stet.**

Fig. 41G, H

Material examined. AD4990: M16923 ([PQ435557](#); 16S: [PQ304663](#)).

Localities. Parrita Seep (1401 m).

Remarks. Genetic data place this specimen within *Pholidoskepia sensu* Kocot et al. 2019. It is possibly a member of *Sandalomeniidae*, pending further molecular characterization of this group.

Mollusca | Solenogastres | Amphimeniidae

***Amphimeniidae* stet.**

Fig. 42A

Material examined. AD4587: M12292 ([PQ449404](#)).

Localities. Mound 12 (996 m).

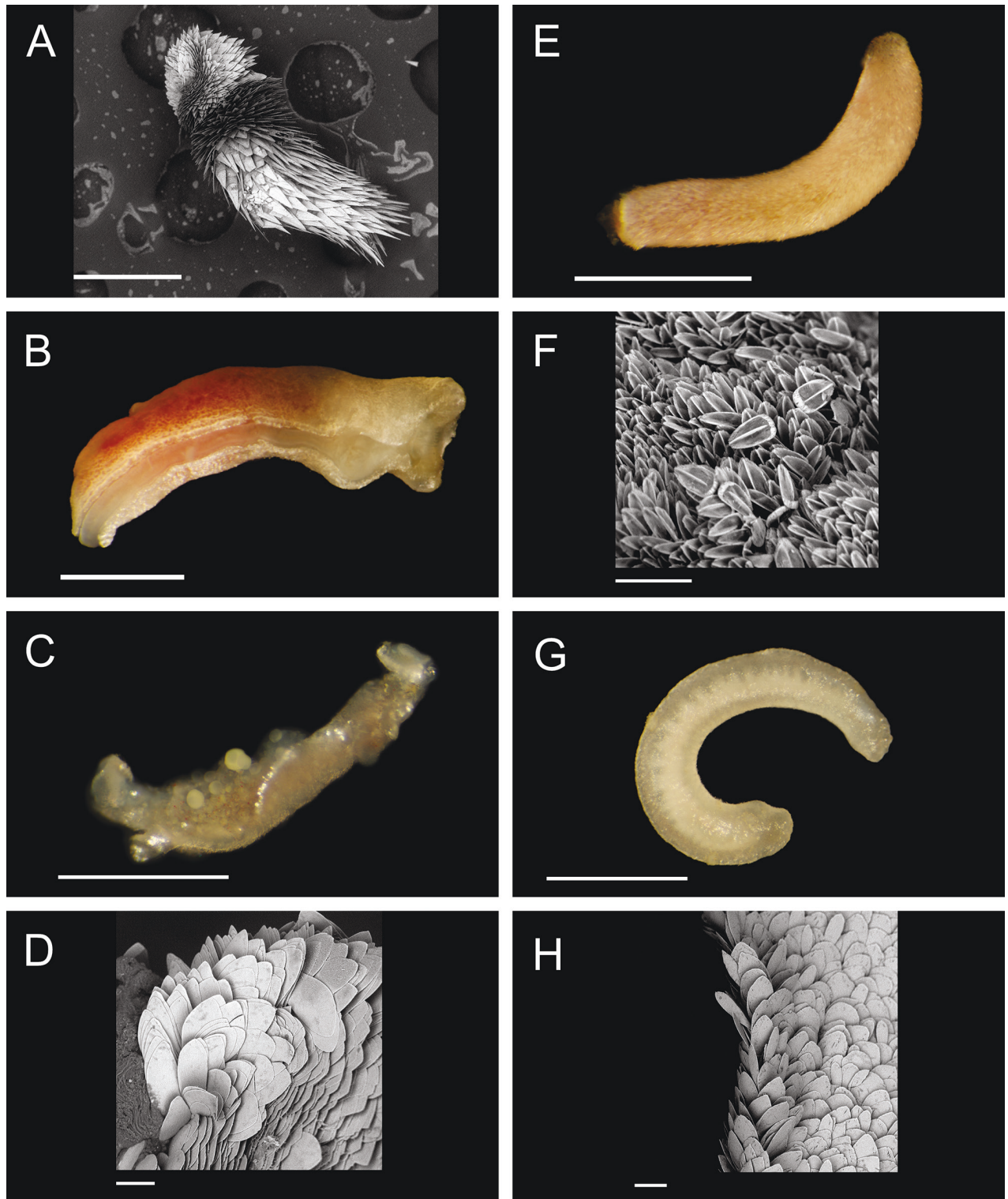


Figure 41. Mollusca: Aplacophora: Caudofoveata and Solenogastres, representative live and SEM images **A** Chaetodermatidae sp. SIO_BIC_M16891 (M16891, SEM) **B** *Neomenia* gen. inc. (M18409) **C** Gymnomeniidae stet. (M16924) **D** Gymnomeniidae stet. (M16924, SEM) **E** *Wirenia* sp. SIO_BIC_M17072 (M17072) **F** *Wirenia* sp. SIO_BIC_M17072 (M17072, SEM) **G** *Pholidoskepia* stet. (M16923) **H** *Pholidoskepia* stet. (M16923, SEM). Scale bars: 0.3 mm (**A**); 1 mm (**B**, **C**, **E**, **G**); 0.02 mm (**D**, **F**, **H**).

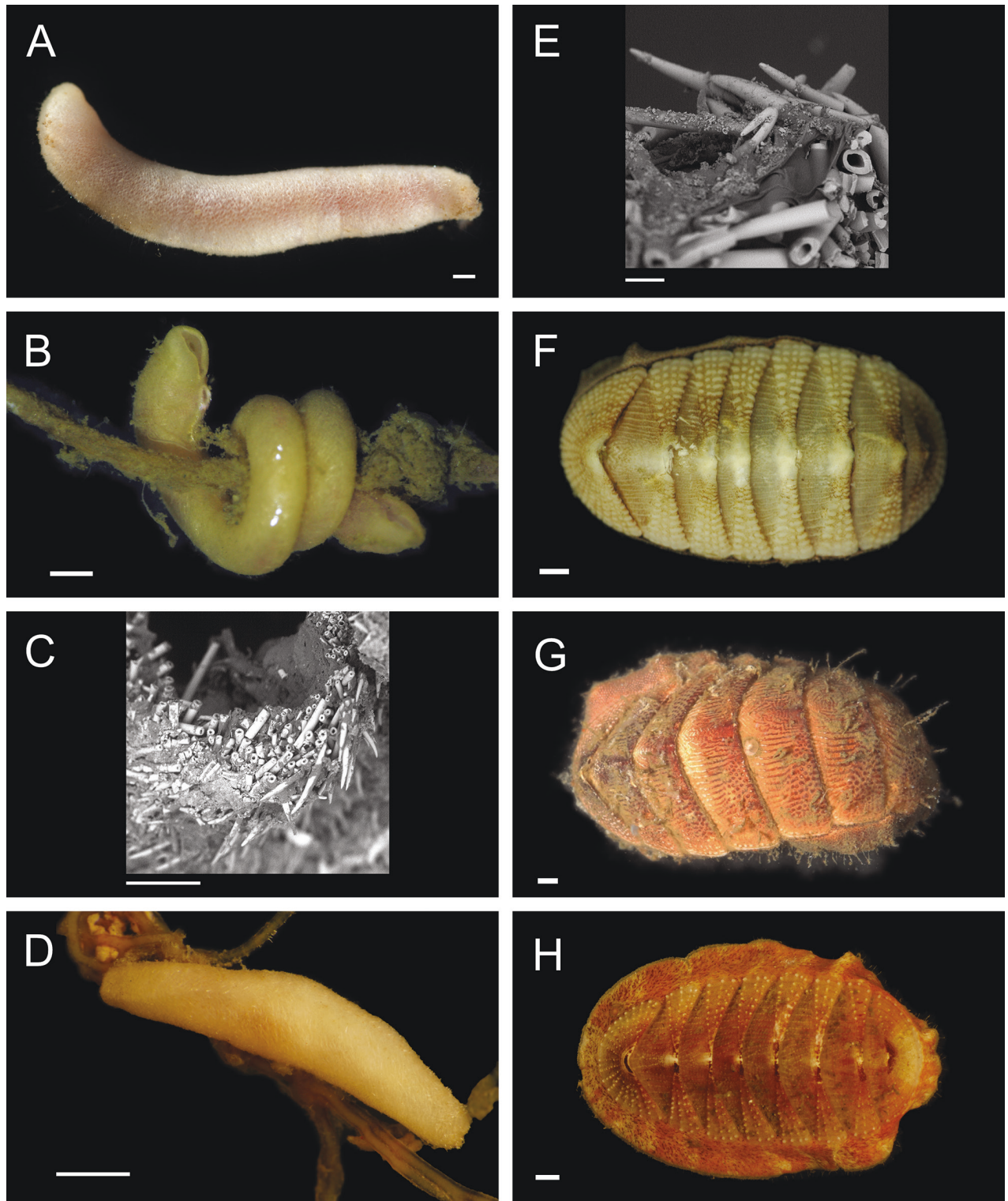


Figure 42. Mollusca: Aplacophora: Solenogastres and Polyplacophora, representative live and SEM images **A** Amphimeniidae stet. (M12292) **B** *Dorymenia* stet. (M11997) **C** *Dorymenia* stet. (M16156, SEM) **D** Pruvotinidae stet. (M16880) **E** Pruvotinidae stet. (M16885, SEM) **F** *Stenosemus* sp. SIO_BIC_M12017 (M12017) **G** *Stenosemus* sp. SIO_BIC_M17044 (M17044) **H** *Tripoplax balaenophila* (M16186). Scale bars: 1 mm (**A, B, D, F–H**); 0.1 mm (**C**); 0.02 mm (**E**).

Mollusca | Solenogastres | Proneomeniidae

***Dorymenia* stet.**

Fig. 42B, C

Material examined. AD4505: M11997; AD4917: M16156 ([PQ435554](#); 16S: [PQ304662](#)).

Localities. Mound 12 (1002 m), Mound 11 (~ 1019–1025 m).

Remarks. M11997 and M16156 are the same morphospecies. M16156 was associated with the octocoral *Swiftia sahlingi* (Co2935).

Mollusca | Solenogastres | Pruvotinidae

Pruvotinidae stet.

Fig. 42D, E

Material examined. AD4978: M16880, M16885 ([PQ435555](#)).

Localities. Mound 12 (997 m).

Remarks. M16880 was associated with a hydroid (Co3639).

Mollusca | Polyplacophora | Chitonida | Ischnochitonidae

***Stenosemus* sp. SIO_BIC_M12017**

Fig. 42F

Material examined. AD4508: M12017 (16S: [PQ304661](#)).

Localities. Parrita Seep (1402 m).

Remarks. An undescribed species.

***Stenosemus* sp. SIO_BIC_M17044**

Fig. 42G

Material examined. S0218: M17044 ([PQ449420](#)).

Localities. Parrita Scar (1364 m).

Remarks. An undescribed species.

***Tripoplax balaenophila* (Schwabe & Sellanes, 2004)**

Fig. 42H

Material examined. AD4512: M12064 ([PQ450384](#); 16S: [PQ304667](#)); AD4918: M16186.

Localities. Quepos Slide (338 m and ~ 344–411 m).

Distribution. Originally described from whale bones at 240 m, within the oxygen minimum zone, off Concepción, central Chile (36°29.9'S, 73°40.8'W) (Schwabe and Sellanes 2004). Specimens identified as *Tripoplax* cf. *balaenophila* have been collected off central Baja California, 530–625 m, in hypoxic conditions near the lower boundary of the oxygen minimum zone (Suárez-Mozo and Hendrickx 2016). Those specimens show morphological variations from the *T. balaenophila* type material and require further taxonomic investigation, including genetic work,

to determine whether they represent a range extension of *T. balaenophila* or a very similar undescribed species (Suárez-Mozo and Hendrickx 2016).

New records. Pending confirmation of the specimens from Mexico, our CRM specimens represent new northern records, new seep records, and a new maximum depth record for this species (specimen M12064, using 344 m as the most conservative value).

Remarks. Consistent with the previous reports of this species in hypoxic conditions, the CRM seep specimens were also collected within the oxygen minimum zone, although not in association with organic falls.

Mollusca | Polyplacophora | Lepidopleurida | Leptochitonidae

Leptochiton is paraphyletic and molecular taxonomic revision is needed (Irisarri et al. 2020).

***Belknapchiton halistreptus* (Dall, 1902)**

Fig. 43A

Material examined. S0219: M17045 ([PQ449421](#)).

Localities. Rio Bongo Scar (609 m).

Distribution. Known only from the type locality off Acapulco, Mexico, 902–3436 m (Dall 1902, 1908; Sirenko et al. 2022). Two subspecies are distinguished by morphology and depth: *B. halistreptus halistreptus* from ~ 3400 m and *B. halistreptus abbreviatus* from ~ 900–1200 m (Dall 1908; Sirenko et al. 2022).

New records. The CRM specimen represents a new southern record and a new minimum depth record for this species.

Remarks. *B. halistreptus* is a member of the recently described genus *Belknapchiton* Sirenko, Saito & Schwabe, 2022. This genus of 22 species includes many of the worldwide chiton specimens obtained from deep water that were formally assigned to *Leptochiton*, and these generally require SEM observations to identify to species. The type species, *B. belknapi* (Dall, 1878), was described from 1840 m off the western Aleutian Islands and is widespread at depths of 100–3724 m in the Pacific, from the Izu-Ogasawara Trench through the Bering Sea and eastern Pacific as far south as central Chile (Sirenko and Sellanes 2016; Sirenko et al. 2022). The CRM specimen has been tentatively identified as one of two similar subspecies of *B. halistreptus*. Both subspecies can be distinguished from *B. belknapi* and other congeners because they have ~ 2× as many gills as *B. belknapi* and a gill row that extends further anterior to approximately the position of the fifth valve (Sirenko et al. 2022).

***Hanleyella* sp. SIO_BIC_M11969**

Fig. 43B

Material examined. AD4501: M11969 (16S: [PQ304660](#)); AD4508: M12015; AD4588: M12131; AD4974: M16766, M16767; AD4978: M16841, M16842, M16883, M16884; AD4987: M16905.

Localities. Mound 12 (~ 1000 m), Parrita Seep (1402 m).

Remarks. These specimens represent an undescribed species with morphological and genetic similarities to *H. oldroydi* (Dall, 1919), which is found from Alaska to Baja California at depths of 18–455 m (Stebbins and Eernisse 2009).

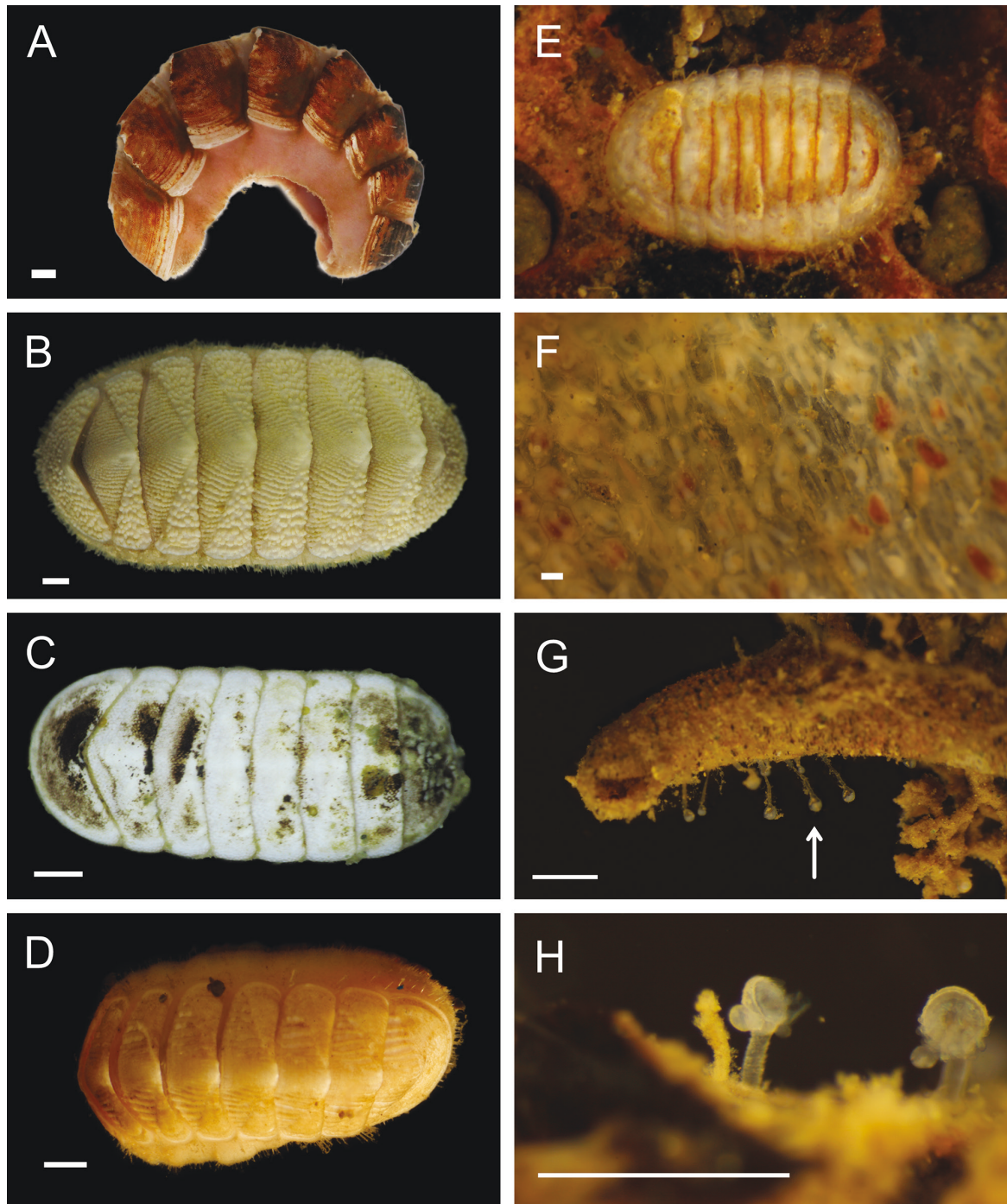


Figure 43. Mollusca: Polyplacophora, Bryozoa, and Entoprocta, representative live images **A** *Belknapchiton halistreptus* (M17045) **B** *Hanleyella* sp. SIO_BIC_M11969 (M11969) **C** *Leptochiton* cf. *americanus* (M12016) **D** *Leptochiton* cf. *incongruus* (M17067) **E** *Leptochiton* sp. SIO_BIC_M17068 (M17068; specimen length estimated < 3 mm, maximum 6 mm) **F** Bryozoa stet. (Ep220) **G** Entoprocta stet. (BI1162) **H** Entoprocta stet. (BI1166, detail). Scale bars: 1 mm.

***Leptochiton* cf. *americanus* Kaas & Van Belle, 1985**

Fig. 43C

Material examined. AD4508: M12016; AD4973: M16725; AD4976: M16814; S0230: M17101, M17104, M17107 ([PQ449425](#)).

Localities. Parrita Seep (1419 m), Jacó Scar (1887 m), Mound Jaguar (1896–2000 m).

Remarks. All specimens except M16725 were associated with naturally occurring or experimentally deployed wood. These specimens may represent an undescribed species or new depth records of *L. americanus*, which was originally described from the Gulf of Panama, 1188 m, and is known from Oregon to Chile, 311–1400 m (Schwabe and Sellanes 2010).

***Leptochiton cf. incongruus* (Dall, 1908)**

Fig. 43D

Material examined. S0219: M17067 ([PQ449422](#)).

Localities. Rio Bongo Scar (661 m).

Remarks. Associated with a naturally occurring wood fall. Possibly an undescribed species or a juvenile of *L. incongruus*, which was described from the Gulf of Panama, 589 m (Dall 1908).

***Leptochiton* sp. SIO_BIC_M17068**

Fig. 43E

Material examined. S0219: M17068 ([PQ449423](#); no voucher remaining after DNA extraction).

Localities. Rio Bongo Scar (661 m).

Remarks. An undescribed species associated with a naturally occurring wood fall.

Bryozoa

Bryozoa stet.

Fig. 43F

Material examined. AD4591: Ep245; AD4924: Ep220.

Localities. Parrita Seep (~ 1400–1410 m), Jacó Scar (~ 1752–1795 m).

Remarks. Encrusting on vesicomysid clams.

Entoprocta

Entoprocta stet.

Fig. 43G, H

Material examined. AD4919: BI1162; AD4921: BI1166.

Localities. Quepos Slide (~ 345–397 m).

Remarks. BI1162 was associated with the tube of a sabellid worm, *Pseudopotamilla* stet. (A8390). BI1166 was associated with a naturally occurring wood fall.

Platyhelminthes

Fecampiida stet.

Fig. 44A, B

Material examined. AD4916: Pt64.

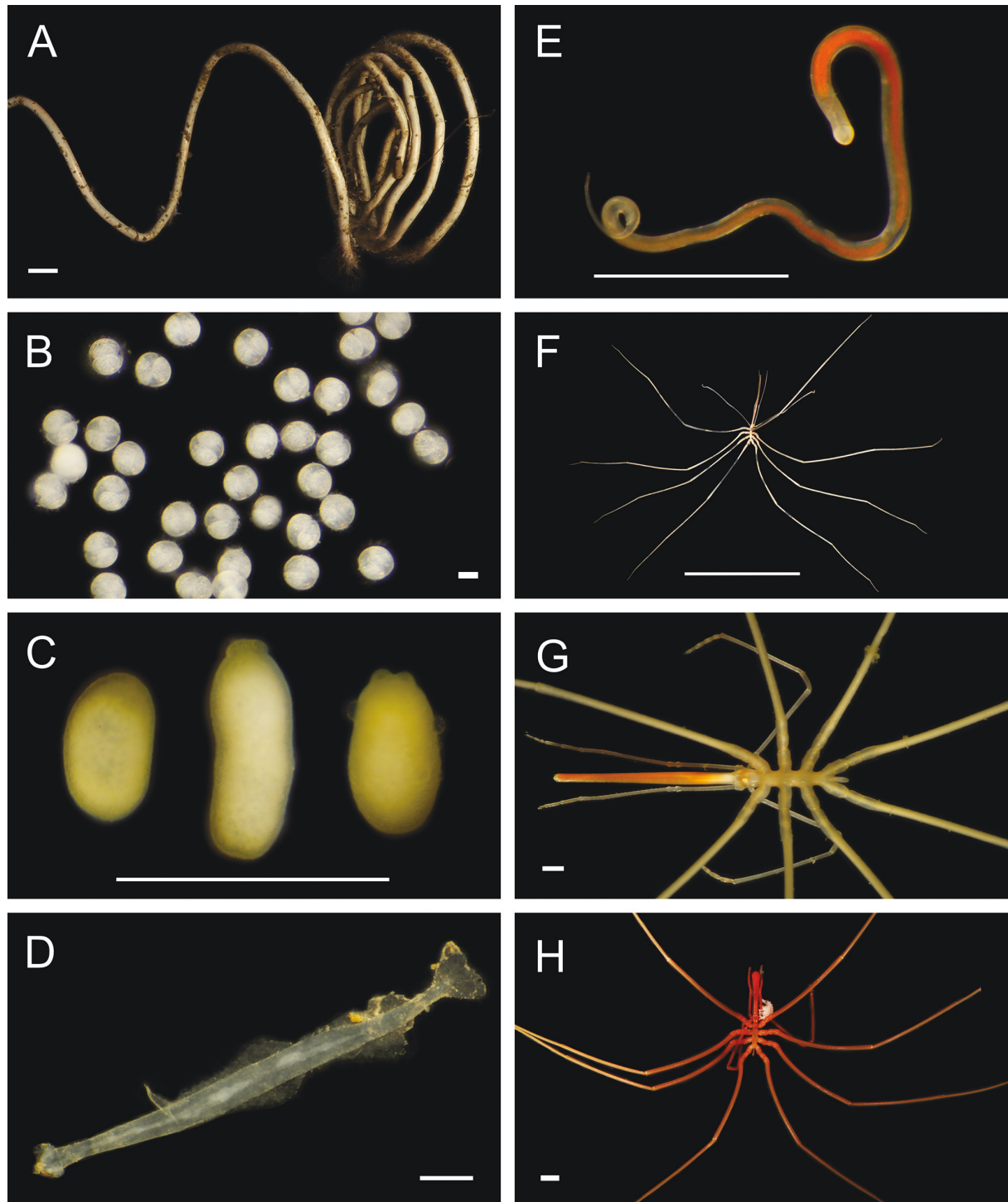


Figure 44. Platyhelminthes, Chaetognatha, Nematoda, and Arthropoda: Pycnogonida, representative live images **A** Fecampiida stet. (Pt64, egg cocoon) **B** Fecampiida stet. (Pt64, detail of eggs) **C** Rhabditophora stet. (Pt72) **D** Chaetognatha stet. (Bl1347) **E** Nematoda stet. (Nto35) **F** *Colossendeis macerrima* (C12792, wide view) **G** *Colossendeis macerrima* (C11151, detail) **H** *Colossendeis* stet. (C13919, wide view). Scale bars: 1 cm (**A**, **H**); 0.1 mm (**B**); 1 mm (**C**–**E**, **G**); 10 cm (**F**).

Localities. Jacó Scar (1854 m).

Remarks. This coiled cocoon of egg capsules was found on soft sediment. Fecampiid cocoons have been previously reported in association with gorgonians in the western Pacific, 92–295 m (Handl and Bouchet 2007).

Rhabditophora stet.

Fig. 44C

Material examined. AD4978: Pt66; AD4985: Pt68; AD4989: Pt72, Pt73.

Localities. Mound 12 (~ 995–1002 m), Jacó Scar (1768 m).

Remarks. Pt72 and Pt73 were associated with a tubeworm bush.

Chaetognatha

Chaetognatha stet.

Fig. 44D

Material examined. AD4985: BI1347.

Localities. Mound 12 (991 m).

Nematoda

Nematoda stet.

Fig. 44E

Material examined. AD4918: Nto35; AD4979: Nto64, Nto65, Nto66, Nto67; S0216: Nto68.

Localities. Quepos Slide (~ 275–400 m).

Arthropoda

We list the major arthropod clades according to the phylogenetic relationships in Regier et al. (2010). See Azofeifa-Solano and Cortés (2020) for a detailed review of deep-sea crustaceans recorded from Costa Rican waters.

Many of the copepod, barnacle, and peracarid morphospecies in this study were represented by small single specimens and may represent undescribed species. To minimize the destruction of diagnostic features, we did not attempt extensive genetic investigation. Specimens are available for loan for future examination.

Arthropoda | Chelicerata | Pycnogonida

We list entries following the phylogeny in Ballesteros et al. (2021). We thank Claudia Arango (Queensland Museum) for assistance with morphological identification of these specimens.

Arthropoda | Chelicerata | Pycnogonida | Colossendeidae

***Colossendeis macerrima* Wilson, 1881**

Fig. 44F, G

Material examined. AD4509: C11151; AD4914: C12792 ([PQ449344](#)).

Localities. Jacó Scar (~ 974–1856 m).

Distribution. Considered cosmopolitan (Munilla and Soler Membrives 2009), originally described from the United States mid-Atlantic coast, 1686 m (Wilson 1881), and previously reported from the Pacific coast of Central America (Hendrickx 2020b).

Remarks. The COI sequence of C12792 was 93.19–95.94% identical to sequences of *Colossendeis macerrima* ([KF603929.1](#), [KF603928.1](#), [JN018213.1](#), [FJ862873.1](#)), with the closest BLASTN matches corresponding to specimens from southern Chile, 510 m (Weis and Melzer 2012). We interpret this 4–7% divergence as intraspecific variation, based on a previous analysis of *C. macerrima* and nine other Chilean pycnogonid species, in which the maximum intra-specific COI divergence was 10.4% and the minimum interspecific COI divergence was 13.36% (Weis and Melzer 2012).

***Colossendeis* stet.**

Figs 44H, 45A

Material examined. S0218: C13919 ([PQ449356](#)).

Localities. Parrita Scar (1153 m).

Remarks. An amphipod, *Mesopleustes abyssorum* (C13920), was attached to the palp of this specimen. The closest COI BLASTN results on GenBank were several species of *Colossendeis* with ~ 88% identity, e.g., *C. colossea* Wilson, 1881 ([FJ716626.1](#), formerly *C. gigas* Hoek, 1881), *C. australis* Hodgson, 1907 ([GQ387003.1](#)), *C. tortipalpis* Gordon, 1932 ([KT202204.1](#)), and *C. macerrima* ([JN018213.1](#)). This level of COI divergence falls between the intraspecific (<10.4%) and interspecific (>13.36%) divergences reported for other pycnogonids (Weis and Melzer 2012), so species-level identification of the CRM specimen will require further investigation.

Arthropoda | Chelicerata | Pycnogonida | Ammotheidae

***Sericosura* sp. SIO_BIC_C13774**

Fig. 45B

Material examined. AD4972: C13774 ([PQ449350](#)).

Localities. Jacó Scar (1795 m).

Remarks. An undescribed species.

***Sericosura* sp. SIO_BIC_C13775**

Fig. 45C

Material examined. AD4972: C13775 ([PQ449351](#)); AD4989: C13865.

Localities. Jacó Scar (1785–1795 m).

Remarks. An undescribed species.

Arthropoda | Chelicerata | Pycnogonida | Phoxichilidiidae

***Anoplodactylus* gen. inc.**

Fig. 45D, E

Material examined. AD4978: C13793 ([PQ449353](#)); AD4985: C13825.

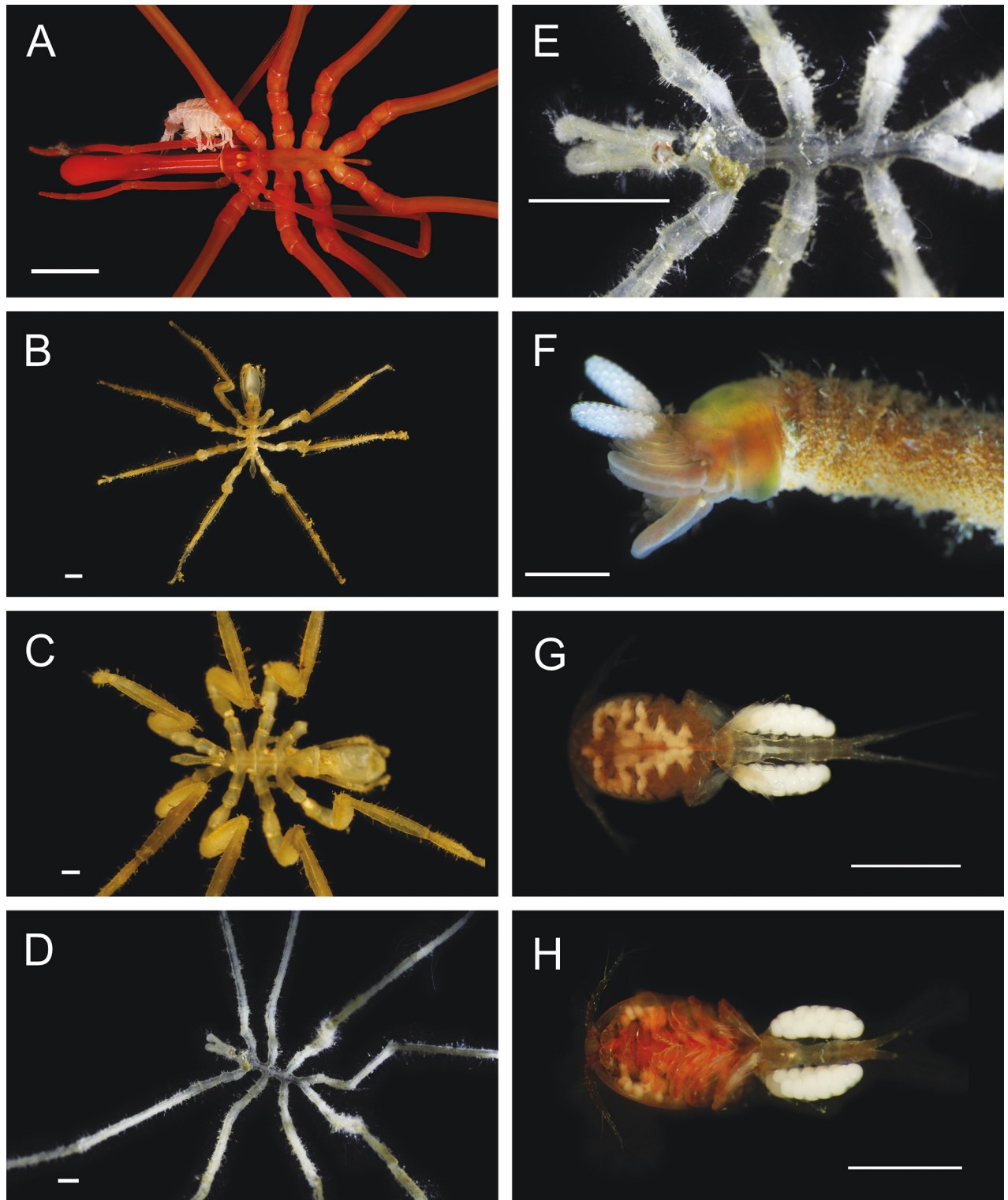


Figure 45. Arthropoda: Pycnogonida and Copepoda, representative live images **A** *Colossendeis* stet. (C13919, detail) **B** *Sericosura* sp. SIO_BIC_C13774 (C13774) **C** *Sericosura* sp. SIO_BIC_C13775 (C13775) **D** *Anoplodactylus* gen. inc. (C13825, wide view) **E** *Anoplodactylus* gen. inc. (C13825, detail) **F** Bradophilidae stet. (A1451) **G** Cyclopoida sp. SIO_BIC_C12780 (C12780, dorsal view) **H** Cyclopoida sp. SIO_BIC_C12780 (C12780, ventral view). Scale bars: 1 cm (**A**); 1 mm (**B–H**).

Localities. Mound 12 (~ 996–1002 m).

Remarks. Most likely *Anoplodactylus*, perhaps an undescribed species (Claudia Arango, pers. comm. 24 July 2022).

Arthropoda | Crustacea | Copepoda

We thank Linsey Sala (Scripps Institution of Oceanography Pelagic Invertebrate Collection) for assistance with these identifications.

Arthropoda | Crustacea | Copepoda | Cyclopoida

Bradophilidae stet.

Fig. 45F

Material examined. AD4511: C14958 (no material remaining); AD4987: C14494.

Localities. Mound 12 (~ 988–1012 m).

Remarks. Egg masses were attached to the flabelligerid *Bradabyssa* cf. *pilosa*: C14958 on host A1451 and C14494 on host A9840.

Cyclopoida sp. SIO_BIC_C12780

Fig. 45G, H

Material examined. AD4503: C11138; AD4587: C11184; AD4910: C12780 ([PQ449343](#)).

Localities. Mound 12 (~ 1000 m).

Remarks. Found in the mantle cavity of the solemyid clam *Acharax* cf. *johnsoni*: C11138 with clam M11980, C12780 with clam M15768.

Cyclopoida sp. SIO_BIC_C12807

Fig. 46A

Material examined. AD4918: C12807.

Localities. Quepos Slide (~ 333–408 m).

Arthropoda | Crustacea | Copepoda | Harpacticoida

Harpacticoida stet.

Fig. 46B

Material examined. AD4988: C13870.

Localities. Mound 11 (~ 1005–1025 m).

Arthropoda | Crustacea | Copepoda | Siphonostomatoida

Caligus stet.

Material examined. AD4503: MZUCR-2770-01 (no images available); AD4505: MZUCR-2771-01 (no images available).

Localities. Mound 11 (~ 1020 m), Mound 12 (~ 1000 m).

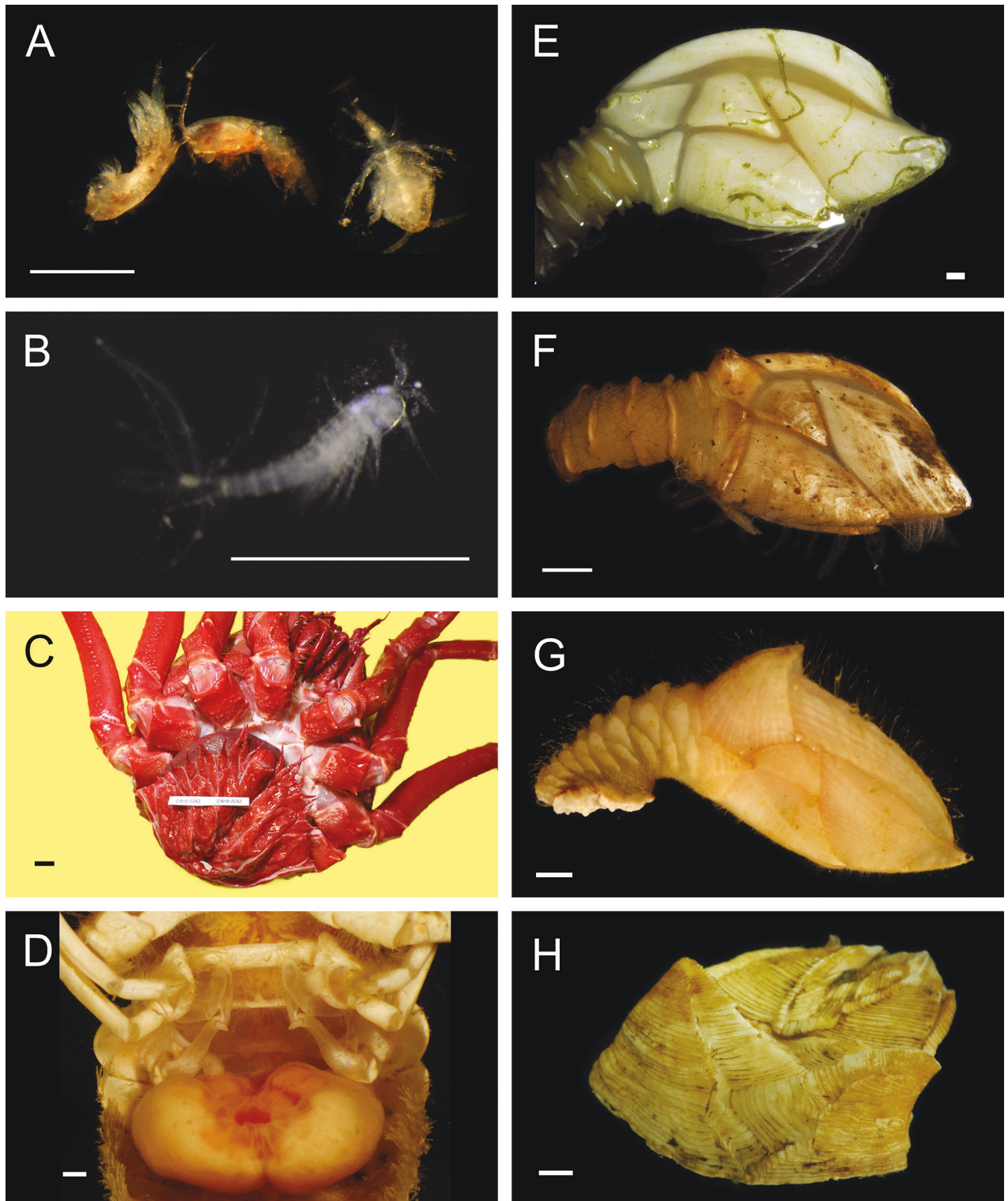


Figure 46. Arthropoda: Copepoda and Cirripedia, representative live images **A** Cyclopoida sp. SIO_BIC_C12807 (C12807) **B** Harpacticoida stet. (C13870) **C** Rhizocephala sp. SIO_BIC_C13776 (C13776) **D** Rhizocephala sp. SIO_BIC_C13875 (C13875) **E** Litoscalpellum sp. SIO_BIC_C11143 (C11143) **F** Litoscalpellum sp. SIO_BIC_C12782 (C12782) **G** Scalpellidae sp. SIO_BIC_C13851 (C13851) **H** Metaverruca gen. inc. (C11148). Scale bars: 1 mm (A, B, D, E, G, H); 1 cm (C, F).

Arthropoda | Crustacea | Thecostraca | Cirripedia

We list entries following the phylogeny in Chan et al. (2021). We thank Hiromi Watanabe (Japan Agency for Marine-Earth Science and Technology) for the morphology-based identifications.

Arthropoda | Crustacea | Thecostraca | Cirripedia | Rhizocephala

Rhizocephala sp. SIO_BIC_C13776

Fig. 46C

Material examined. AD4975: C13776 ([PQ448998](#)).

Localities. Mound 12 (1000 m).

Remarks. Parasite of a lithodid crab, *Lithodes panamensis* (C13787). The closest COI BLASTN results on GenBank were within Peltogastridae: *Briarosaccus* sp. ([OR466125.1](#), 84.23% identity), *Peltogaster boschmai* Reinhard, 1944 from the San Juan Islands, Washington, USA ([MN138416.1](#), 80.00% identity), and several sequences of *P. lineata* Shiino, 1943 from Korea and Japan (e.g., [MK604142.1](#), 78.82% identity).

Rhizocephala sp. SIO_BIC_C13875

Fig. 46D

Material examined. AD4989: C13875 ([PQ449355](#)).

Localities. Jacó Scar (1762 m).

Remarks. Parasite of a squat lobster, *Munidopsis alvisca* (C13876). The closest COI BLASTN results on GenBank were several species of *Lernaeodiscus* (Peltogastridae), e.g., *L. ingolfi* Boschma, 1928 from Norway ([MN605966.1](#), 80.62% identity) and *L. rybakovi* Korn, Golubinskaya, Rees, Glenner & Høeg, 2020 from Vostok Bay, Russia ([MN605964.1](#), 78.81% identity).

Arthropoda | Crustacea | Thecostraca | Cirripedia | Thoracica | Scalpellomorpha

Litoscalpellum sp. SIO_BIC_C11143

Fig. 46E

Material examined. AD4504: C11143.

Localities. Mound 11 (~ 1004–1011 m).

Remarks. Attached to a vestimentiferan tubeworm. *Litoscalpellum* is polyphyletic and revision is required (Linse et al. 2013).

Litoscalpellum sp. SIO_BIC_C12782

Fig. 46F

Material examined. AD4913: C12782.

Localities. Jacó Scar (1885 m).

Scalpellidae sp. SIO_BIC_C13851

Fig. 46G

Material examined. AD4990: C13851.

Localities. Parrita Seep (1401 m).

Arthropoda | Crustacea | Thecostraca | Cirripedia | Thoracica |
Verrucomorpha

***Metaverruca* gen. inc.**

Fig. 46H

Material examined. AD4508: C11148.

Localities. Parrita Seep (~ 1401–1419 m).

Remarks. Likely *Metaverruca*. Attached to a tubeworm, *Lamellibrachia barhami*.

***Newmaniverruca* gen. inc.**

Fig. 47A

Material examined. AD4916: C12794.

Localities. Jacó Scar (1611 m).

Remarks. Likely *Newmaniverruca*.

Verrucidae sp. SIO_BIC_C11144

Fig. 47B

Material examined. AD4506: C11144.

Localities. Parrita Seep (~ 1030–1179 m).

Remarks. Likely *Altiverruca* or *Newmaniverruca*.

Arthropoda | Crustacea | Thecostraca | Cirripedia | Thoracica |
Balanomorpha

Pyrgomatidae stet.

Fig. 47C

Material examined. AD4923: C12815.

Localities. Parrita Seep (~ 1041–1094 m).

Remarks. Associated with a coralliid (Co2947).

Arthropoda | Crustacea | Malacostraca | Hoplocarida | Stomatopoda |
Squillidae

***Squilla biformis* Bigelow, 1891**

Fig. 47D, E

Reference. Koga and Rouse (2021) for the mitochondrial genome and phylogenetic analysis of C13808.

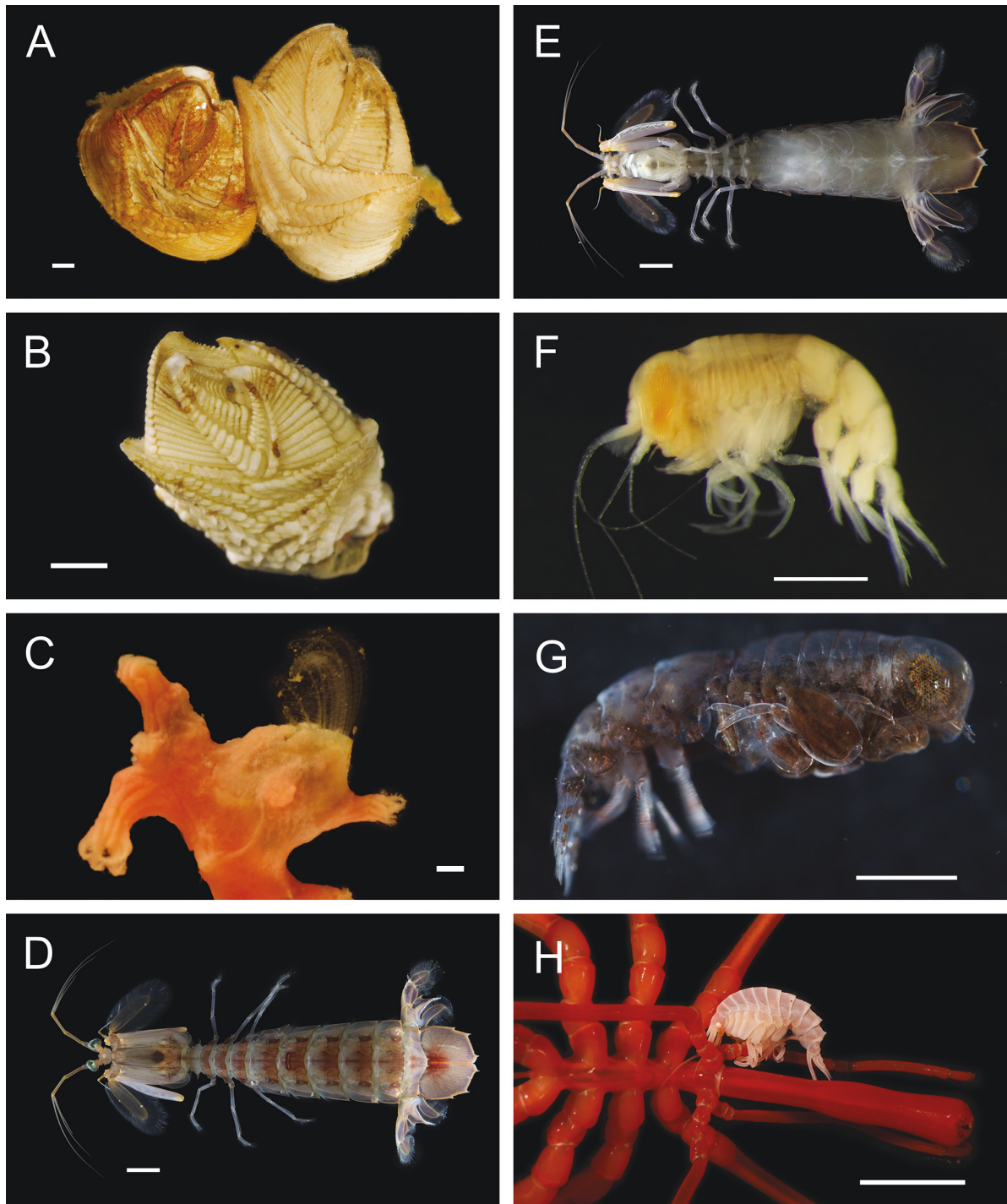


Figure 47. Arthropoda: Cirripedia, Stomatopoda, and Amphipoda, representative images. Live specimens are depicted unless otherwise specified **A** *Newmaniverruca* gen. inc. (C12794) **B** Verrucidae sp. SIO_BIC_C11144 (C11144) **C** Pyrgomatidae stet. (C12815) **D** *Squilla biformis* (C13807, dorsal view) **E** *Squilla biformis* (C13807, ventral view) **F** Hyperiidea stet. (C15409, preserved specimen) **G** *Lycaea pulex* (C14397) **H** *Mesopleustes abyssorum* (C13920). Scale bars: 1 mm (**A–C, F, G**); 1 cm (**D, E, H**).

Material examined. AD4979: C13807 ([PQ449354](#)), C13808 ([MW867305](#)); AD4986: MZUCR-3732-01.

Localities. Quepos Slide (~ 380–395 m).

Distribution. Originally described from La Paz, Gulf of California, at 205 m (Bigelow 1894) and distributed south to Peru at depths of 28–518 m (Hendrickx

and Salgado-Barragán 1991; Hendrickx and López 2020). *S. biformis* has been studied from Pacific Costa Rica at depths of 131–350 m (Camp and Kuck 1990; Wehrtmann and Echeverría-Sáenz 2007; Hernáez et al. 2011; Azofeifa-Solano and Cortés 2020).

Remarks. Telson morphology indicates that both specimens are female. To our knowledge this study is the first report of this well-known local species in the vicinity of seeps, notably within the oxygen minimum zone as characterized by Levin et al. (2015).

Arthropoda | Crustacea | Malacostraca | Peracarida

Several available checklists of eastern Pacific deep-sea peracarids (Hendrickx 2020a) may be informative for future work on these specimens. Unless otherwise stated, we list entries following the phylogenies in Schwentner et al. (2018) and Höpel et al. (2022).

Arthropoda | Crustacea | Malacostraca | Peracarida | Lophogastrida

Eucopia sculpticauda Faxon, 1893

Material examined. AD4513: MZUCR-2818-01 (no image available).

Localities. Jacó Scar (~ 1800 m).

Distribution. Originally described from several stations in the Gulf of Panama and off the Galápagos Islands, 1618–2487 m (Faxon 1893, 1895). Reported with a wide latitudinal distribution in the Indo-Pacific and Atlantic Oceans, from ~ 1000–7526 m depth (Kou et al. 2019).

Arthropoda | Crustacea | Malacostraca | Peracarida | Amphipoda

We list entries following the World Amphipoda Database (Horton et al. 2022), acknowledging the need for further systematic work as discussed in the molecular phylogeny of Copilaş-Ciocianu et al. (2020).

Arthropoda | ... | Amphipoda | Hyperidei

We thank Linsey Sala for these identifications.

Hyperidei

Fig. 47F

Material examined. AT15-59 Plankton Tow 6: C15409.

Localities. Jacó Summit (~ 350 m depth, ~ 400 m above the seafloor).

Lycaea pulex Marion, 1874

Fig. 47G

Material examined. AT15-59 Plankton Tow 6: C14397.

Localities. Jacó Summit (~ 350 m depth, ~ 400 m above the seafloor).

Distribution. Considered common and widespread in tropical to warm-temperate oceans worldwide, typically 0–500 m depth (Zeidler 2021).

Arthropoda | ... | Amphipoda | Amphilochidea | Amphilochida |
Amphilochidira | Amphilochoidea | Pleustidae

***Mesopleustes abyssorum* (Stebbing, 1888)**

Fig. 47H

Material examined. S0218: C13920.

Localities. Parrita Scar (1153 m).

Distribution. Originally described from the subantarctic Indian Ocean off South Africa, 2926 m (Stebbing 1888), *M. abyssorum* is considered “probably cosmopolitan” at depths below ~ 700 m (Barnard 1967). In the eastern Pacific, *M. abyssorum* has been reported from 3479 m off Baja California (Barnard 1967).

Remarks. Observed *in situ* attached to the palp of a pycnogonid, *Colossendeis macerrima* (C13919) and remained attached after collection.

***Stenopleustes* gen. inc.**

Fig. 48A

Material examined. AD4507: C11146 ([PQ450405](#)), C14496; AD4916: C12788; AD4922: C12811.

Localities. Mound 12 (~ 967 m), Jacó Scar (~ 1852–1855 m), Parrita Scar (~ 1659–1667 m).

Remarks. C12811 was associated with an antipatharian coral (specific host not recorded). This morphospecies was identified as most likely *Stenopleustes* (Pleustidae), but an alternative identification is Stenothoidae. More detailed morphological examination of vouchers is needed.

Arthropoda | ... | Amphipoda | Amphilochidea | Amphilochida |
Amphilochidira | Amphilochoidea | Sebidae

***Seba* stet.**

Fig. 48B

Material examined. S0230: C13970.

Localities. Mound Jaguar (1896 m).

Remarks. Associated with a naturally occurring wood fall.

Arthropoda | ... | Amphipoda | Amphilochidea | Amphilochida |
Amphilochidira | Amphilochoidea | Stenothoidae

***Stenothoe* sp. SIO_BIC_C13857**

Fig. 48C

Material examined. AD4985: C13857.

Localities. Mound 12 (991 m).

Remarks. This morphospecies lacks eyes.

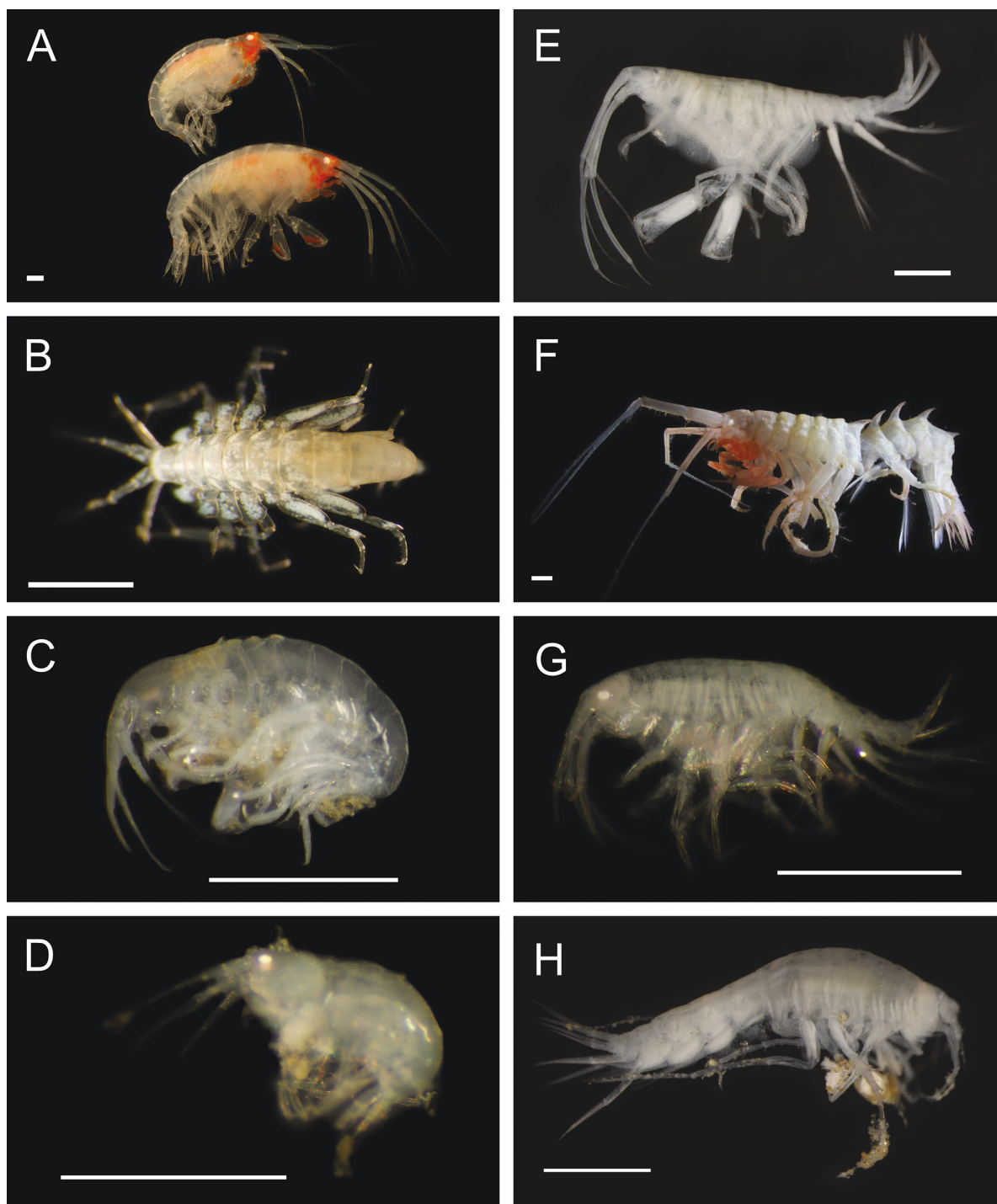


Figure 48. Arthropoda: Amphipoda, representative images. Live specimens are depicted unless otherwise specified **A** *Stenopleustes* gen. inc. (C12811) **B** *Seba* stet. (C13970) **C** *Stenothoe* sp. SIO_BIC_C13857 (C13857) **D** *Stenothoe* sp. SIO_BIC_C13867 (C13867) **E** *Stenula* stet. (C13784, preserved specimen) **F** *Rhachotropis* stet. (C13798) **G** *Idunella* gen. inc. (C13856) **H** *Monoculodes* stet. (C12801, preserved specimen). Scale bars: 1 mm.

***Stenothoe* sp. SIO_BIC_C13867**

Fig. 48D

Material examined. AD4987: C13867.

Localities. Mound 12 (999 m).

Remarks. This morphospecies has eyes.

Stenula stet.

Fig. 48E

Material examined. AD4974: C13784.

Localities. Mound 12 (992 m).

Remarks. Associated with experimental deployments of bone and wood.

Arthropoda | ... | Amphipoda | Amphilochidea | Amphilochida | Eusirida |
Eusiroidea | Eusiridae

Rhachotropis stet.

Fig. 48F

Material examined. AD4507: C14497; AD4976: C13798.

Localities. Jacó Scar (1887 m), Parrita Scar (~ 1659–1667 m).

Remarks. Specimen C13798 may have been associated with experimentally deployed wood.

Arthropoda | ... | Amphipoda | Amphilochidea | Amphilochida | Eusirida |
Liljeborgioidea | Liljeborgiidae

Idunella gen. inc.

Fig. 48G

Material examined. AD4985: C13856.

Localities. Mound 12 (991 m).

Remarks. This morphospecies has eyes. It is most likely *Idunella* (Liljeborgiidae), but an alternative identification is *Stenopleustes* (Pleustidae). Further morphological examination is needed.

Arthropoda | ... | Amphipoda | Amphilochidea | Amphilochida |
Oedicerotidira | Oedicerotoidea | Oedicerotidae

Monoculodes stet.

Fig. 48H

Material examined. AD4507: C14495; AD4589: C11188 ([PQ449341](#)); AD4917: C12785; AD4922: C12801.

Localities. Mound 12 (965–997 m), Parrita Scar (~ 1659–1667 m).

Remarks. C12785 was associated with the antipatharian coral *Lillipathes ritamariae*. C12801 was associated with a basket star, *Gorgonocephalus stet.* (E7064).

Arthropoda | ... | Amphipoda | Amphilochidea | Lysianassida |
Haustoriidira | Haustorioidea | Phoxocephalidae

Phoxocephalinae subfam. inc.

Fig. 49A

Material examined. AD4984: C13858.

Localities. Mound 12 (998 m).

Arthropoda | ... | Amphipoda | Amphilochidea | Lysianassida |
Lysianassidira | Aristioidea | Ambasiidae

***Ambasiella* stet.**

Fig. 49B

Material examined. S0218: C13917.

Localities. Parrita Scar (1988 m).

Remarks. Associated with a xenophyophore.

Arthropoda | ... | Amphipoda | Amphilochidea | Lysianassida |
Lysianassidira | Lysianassoidea | Tryphosidae

***Orchomene* stet.**

Fig. 49C

Material examined. S0230: C13972, C13977.

Localities. Mound Jaguar (1895–1909 m).

Tryphosidae stet.

Fig. 49D

Material examined. AD4974: C13785; AD4985: C13854.

Localities. Mound 12 (992–1002 m).

Remarks. C13785 was associated with experimental deployments of bone and wood.

Arthropoda | ... | Amphipoda | Amphilochidea | Lysianassida |
Lysianassidira | Lysianassoidea incertae sedis

***Ambasiopsis* stet.**

Fig. 49E

Material examined. AD4972: C13789, C13790.

Localities. Jacó Scar (1845 m).

Remarks. Associated with experimentally deployed pig bones.

Arthropoda | ... | Amphipoda | Amphilochidea | Lysianassida |
Lysianassidira | Stegocephaloidea | Stegocephalidae

Stegocephalidae stet.

Fig. 49F

Material examined. S0230: C13976, C13978.

Localities. Mound Jaguar (1895–1908 m).

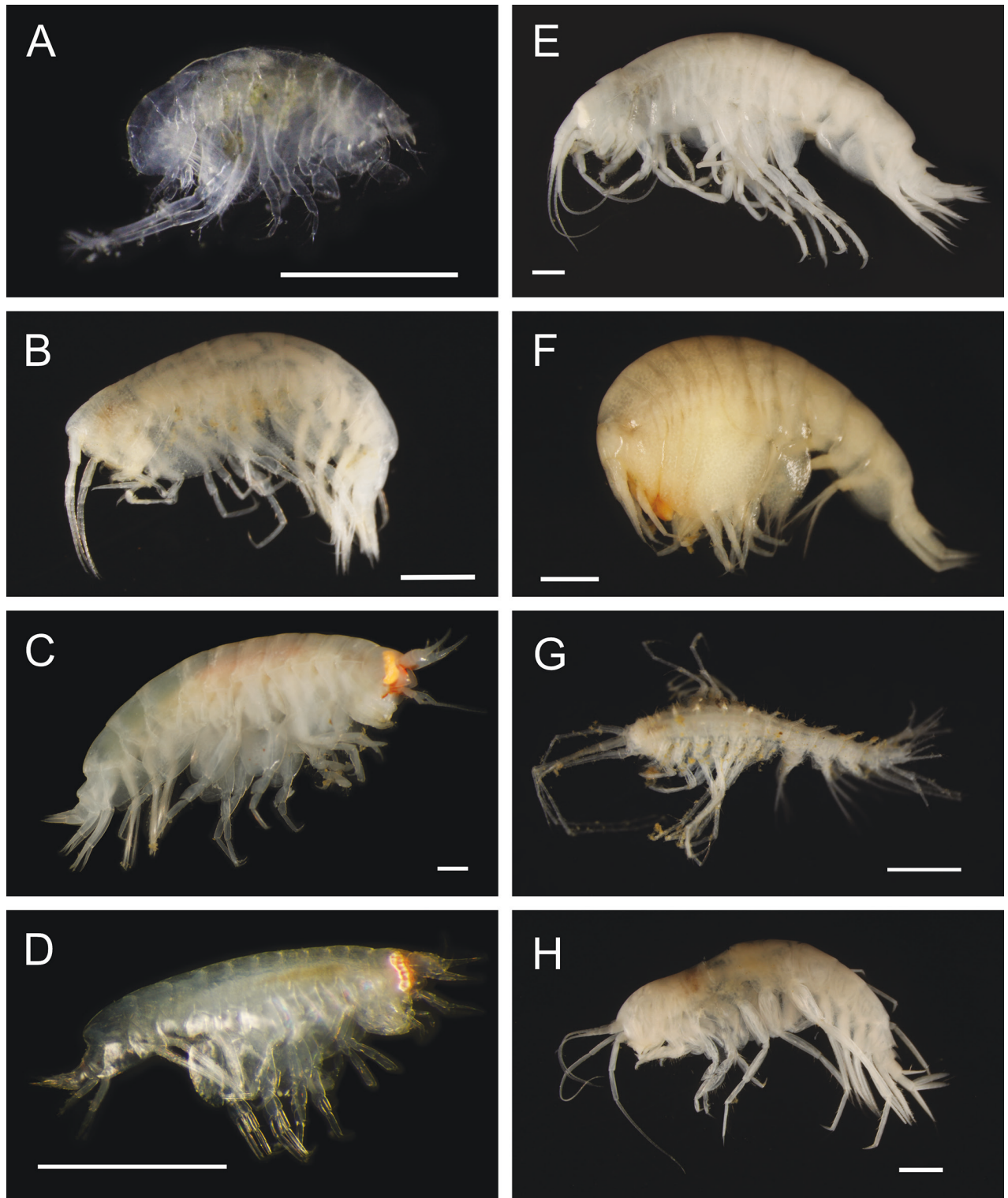


Figure 49. Arthropoda: Amphipoda, representative images. Live specimens are depicted unless otherwise specified **A** Phoxocephalinae subfam. inc. (C13858) **B** *Ambasiella* stet. (C13917, preserved specimen) **C** *Orchomene* stet. (C13972) **D** Tryphosidae stet. (C13854) **E** *Ambasiopsis* stet. (C13790, preserved specimen) **F** Stegocephalidae stet. (C13976, preserved specimen) **G** *Lepechinella* stet. (C13929, preserved specimen) **H** *Pardalisca* stet. (C13928, preserved specimen). Scale bars: 1 mm.

Arthropoda | ... | Amphipoda | Amphilochidea | Lysianassida |
Synopiidira | Dexaminoidea | Lepechinellidae

***Lepechinella* stet.**

Fig. 49G

Material examined. S0219: C13929.

Localities. Rio Bongo Scar (~ 480–650 m).

Arthropoda | ... | Amphipoda | Amphilochidea | Lysianassida |
Synopiidira | Dexaminoidea | Pardaliscidae

***Pardalisca* stet.**

Fig. 49H

Material examined. S0219: C13928.

Localities. Rio Bongo Scar (661 m).

Remarks. Associated with a naturally occurring wood fall.

Arthropoda | ... | Amphipoda | Amphilochidea | Lysianassida |
Synopiidira | Synopioidea | Argissidae

***Argissa* sp. SIO_BIC_C13930**

Fig. 50A

Material examined. S0219: C13930.

Localities. Rio Bongo Scar (~ 480–650 m).

Remarks. Possibly an undescribed species, requiring further comparison. *Argissa* is currently accepted as monotypic, with *Argissa hamatipes* (Norman, 1869) reportedly occurring across the northern hemisphere at depths of 4–1096 m (Barnard 1967; Winfield et al. 2020). A specimen provisionally identified as *A. hamatipes* has been recorded from western Mexico, minimum depth 1720 m (Barnard 1967).

Arthropoda | ... | Amphipoda | Senticaudata | Corophiida | Caprellidira |
Photoidea | Ischyroceridae

***Bonnierella* stet.**

Fig. 50B

Material examined. AD4916: C12789; AD4972: C13797; AD4974: C13786.

Localities. Mound 12 (992 m), Jacó Scar (~ 1646–1751 m).

Remarks. C13786 was associated with experimental deployments of bone and wood.

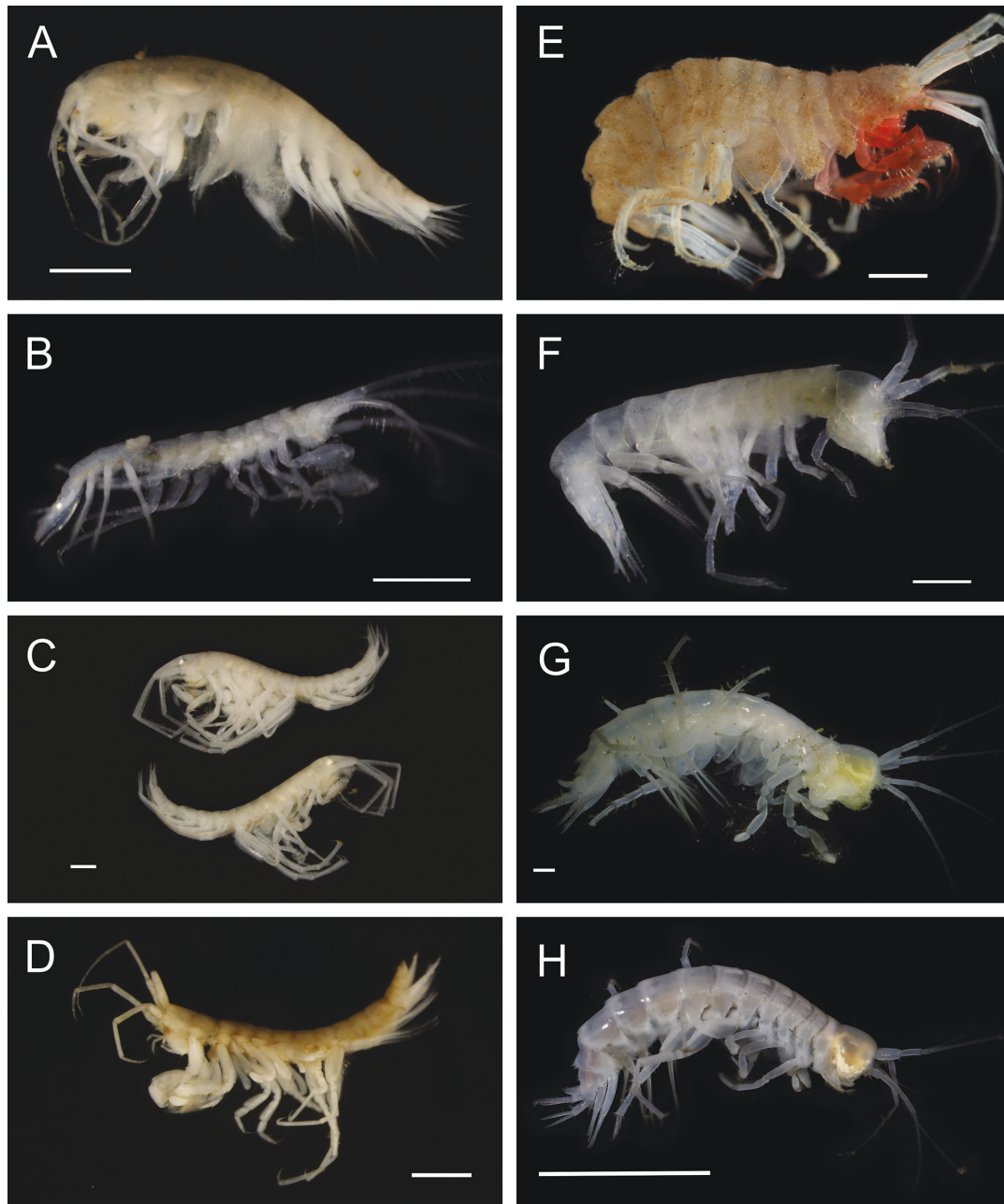


Figure 50. Arthropoda: Amphipoda, representative images. Live specimens are depicted unless otherwise specified **A** *Argissa* sp. SIO_BIC_C13930 (C13930, preserved specimen) **B** *Bonnierella* stet. (C13797) **C** *Gammaropsis* gen. inc. (C13963, preserved specimen) **D** *Bemlos* gen. inc. (C13931, preserved specimen) **E** Protomeleiinae stet. (C14396) **F** *Oradarea* stet. (C13796) **G** *Abludomelita* gen. inc. (C11141) **H** *Abludomelita* stet. (C13896). Scale bars: 1 mm (**A–G**); 1 cm (**H**).

Arthropoda | ... | Amphipoda | Senticaudata | Corophiida | Caprellidira |
Photoidea | Photidae

***Gammaropsis* gen. inc.**

Fig. 50C

Material examined. S0230: C13963.

Localities. Mound Jaguar (1895 m).

Remarks. One specimen is a brooding female. Further morphological examination is needed for a more definitive identification.

Arthropoda | ... | Amphipoda | Senticaudata | Corophiida | Corophiidira |
Aoroidea | Aoridae

***Bemlos* gen. inc.**

Fig. 50D

Material examined. S0219: C13931.

Localities. Rio Bongo Scar (~ 480–650 m).

Arthropoda | ... | Amphipoda | Senticaudata | Corophiida | Corophiidira |
Corophioidea | Corophiidae

***Protomedeiinae* stet.**

Fig. 50E

Material examined. AT15-59 MC4: C14396.

Localities. Near Mound 11 (1031 m).

Remarks. This specimen was collected in a sediment core adjacent to Mound 11, ca 400 m from known sites of active seepage and likely representing the far-transition zone to the surrounding environment.

Arthropoda | ... | Amphipoda | Senticaudata | Hadziida | Hadziidira |
Calliopioida | Calliopiidae

***Oradarea* stet.**

Fig. 50F

Material examined. AD4913: C12784; AD4922: C12800; AD4923: C14500; AD4976: C13796; AD4988: C13871; S0230: C13971, C13979.

Localities. Mound 12 (1001 m), Mound 11 (1010 m), Parrita Seep (~ 1037–1108 m), Jacó Scar (~ 1800–1900 m), Mound Jaguar (1896 m).

Remarks. Several specimens (C13871, C13971, C13979; possibly also C12784 and C13796) were associated with naturally occurring wood falls.

Arthropoda | ... | Amphipoda | Senticaudata | Hadziida | Hadziidira |
Hadzioidea | Melitidae

***Abludomelita* gen. inc.**

Fig. 50G

Material examined. AD4503: C11141; AD4923: C14498, C14499.

Localities. Mound 12 (990 m), Parrita Seep (~ 1037–1108 m).

Remarks. Specimen C11141 was associated with a naturally occurring wood fall. Specimens C14498 and C14499 were likely associated with corals.

***Abludomelita* stet.**

Fig. 50H

Material examined. AD4972: C13791; S0212: C13896, MZUCR-3756-01; S0230: C13975, C13980.

Localities. Jacó Scar (1845–1896 m), Mound Jaguar (1896–2000 m).

Remarks. Associated with experimentally deployed pig bones (C13791) or with naturally occurring wood falls (C13896, C13975, C13980).

Arthropoda | Crustacea | Malacostraca | Peracarida | Tanaidacea |
Apseudomorpha | Apseudoidea | Apseudidae

***Apseudidae* stet.**

Fig. 51A

Material examined. S0230: C13973.

Localities. Mound Jaguar (1909 m).

Arthropoda | Crustacea | Malacostraca | Peracarida | Tanaidacea |
Tanaidomorpha | Paratanaoidea | Agathotanaidae

***Agathotana* gen. inc.**

Fig. 51B

Material examined. AD4979: C13815.

Localities. Quepos Slide (397 m).

Remarks. Collected 2 cm deep in a sediment core. Identification is uncertain because the specimen is a juvenile.

***Paranarthrura* stet.**

Fig. 51C

Material examined. AD4989: C13874.

Localities. Jacó Scar (1768 m).

Remarks. Collected from a vestimentiferan tubeworm bush.

Arthropoda | Crustacea | Malacostraca | Peracarida | Tanaidacea |
Tanaidomorpha | Paratanaoidea | Anarthruridae

***Anarthrurella* gen. inc.**

Fig. 51D

Material examined. AD4971: C13777.

Localities. Jacó Scar (1824 m).

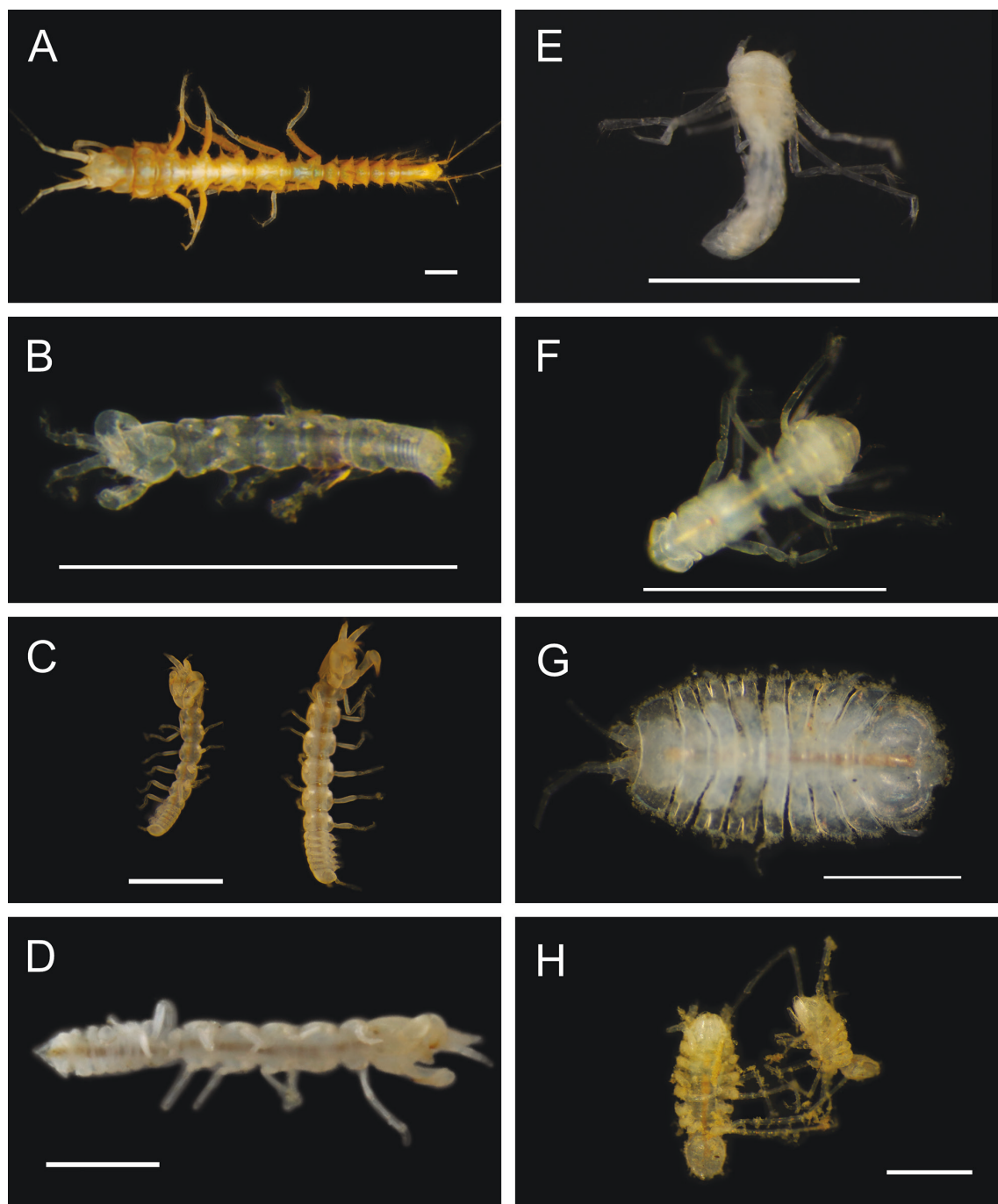


Figure 51. Arthropoda: Tanaidacea and Isopoda, representative images. Live specimens are depicted unless otherwise specified **A** Apseudidae stet. (C13973) **B** Agathotana gen. inc. (C13815) **C** Paranarthra stet. (C13874) **D** Anarthrella gen. inc. (C13777) **E** Desmosoma stet. (C13855, preserved specimen) **F** Mirabilicoxa stet. (C13866) **G** Janthura stet. (C13848) **H** Munna stet. (C13849). Scale bars: 1 mm.

Arthropoda | Crustacea | Malacostraca | Peracarida | Isopoda | Asellota
| Janiroidea | Desmosomatidae

***Desmosoma* stet.**

Fig. 51E

Material examined. AD4985: C13855.

Localities. Mound 12 (1002 m).

Mirabilicoxa stet.

Fig. 51F

Material examined. AD4987: C13866.

Localities. Mound 12 (999 m).

Arthropoda | Crustacea | Malacostraca | Peracarida | Isopoda | Asellota
| Janiroidea | Janiridae

Janthura stet.

Fig. 51G

Material examined. AD4988: C13848.

Localities. Mound 11 (1010 m).

Remarks. Associated with a naturally occurring wood fall.

Arthropoda | Crustacea | Malacostraca | Peracarida | Isopoda | Asellota
| Janiroidea | Munnidae

Munna gen. inc.

Fig. 51H

Material examined. AD4988: C13872.

Localities. Mound 11 (1010 m).

Remarks. Associated with a naturally occurring wood fall.

Munna stet.

Fig. 52A

Material examined. AD4988: C13849.

Localities. Mound 11 (1010 m).

Remarks. Associated with a naturally occurring wood fall.

Arthropoda | Crustacea | Malacostraca | Peracarida | Isopoda | Asellota
| Janiroidea | Munnopsidae

Betamorpha stet.

Fig. 52B

Material examined. S0230: C13974.

Localities. Mound Jaguar (1909 m).

Eurycope stet.

Fig. 52C

Material examined. AD4990: C13850.

Localities. Parrita Seep (1401 m).

Arthropoda | Crustacea | Malacostraca | Peracarida | Isopoda | Asellota
| Stenetrioidea | Stenetriidae

***Tenupedunculus* stet.**

Fig. 52D

Material examined. AD4987: C13823.

Localities. Mound 12 (999 m).

Remarks. Associated with sediment samples.

Arthropoda | Crustacea | Malacostraca | Peracarida | Isopoda |
Cymothoida | Cymothooidea | Gnathiidae

Gnathiidae stet.

Fig. 52E

Material examined. AD4978: C13801.

Localities. Mound 12 (~ 996–999 m).

Arthropoda | Crustacea | Malacostraca | Peracarida | Cumacea

***Paraleucon* sp. SIO_BIC_C11152**

Fig. 52F

Material examined. AD4510: C11152; AD4987: C13860; AD4988: C13873.

Localities. Jacó Summit (741 m), Mound 11 (1010 m), Mound 12 (1010 m).

Remarks. Likely an undescribed species. C13860 was associated with a bed of small clams. C13873 was associated with a naturally occurring wood fall.

Arthropoda | Crustacea | Malacostraca | Eucarida | Decapoda

We list entries following the phylogeny in Wolfe et al. (2019).

Arthropoda | ... | Decapoda | Pleocyemata | Stenopodidea |
Spongicolidae

***Spongicoloides galapagensis* Goy, 1980**

Fig. 52G, H

Material examined. AD4923: C12735, C12814 ([PQ449346](#), [PQ449347](#)), C12820.

Localities. Parrita Seep (~ 1091–1098 m).

Distribution. Described from 717 m off the Galápagos Islands (Goy 1980). Comparison to records of *S. aff. galapagensis* associated with euplectellid sponges at seeps at ~ 1000 m off central Chile (Guzmán and Sellanes 2011) would be informative.

New records. The CRM specimens represent new northern records and new maximum depth records for this species (1091 m as the most conservative value).

Remarks. Associated with a euplectellid sponge, *Dictyaulus* gen. inc. (P1687). One specimen of C12820 is an ovigerous female.

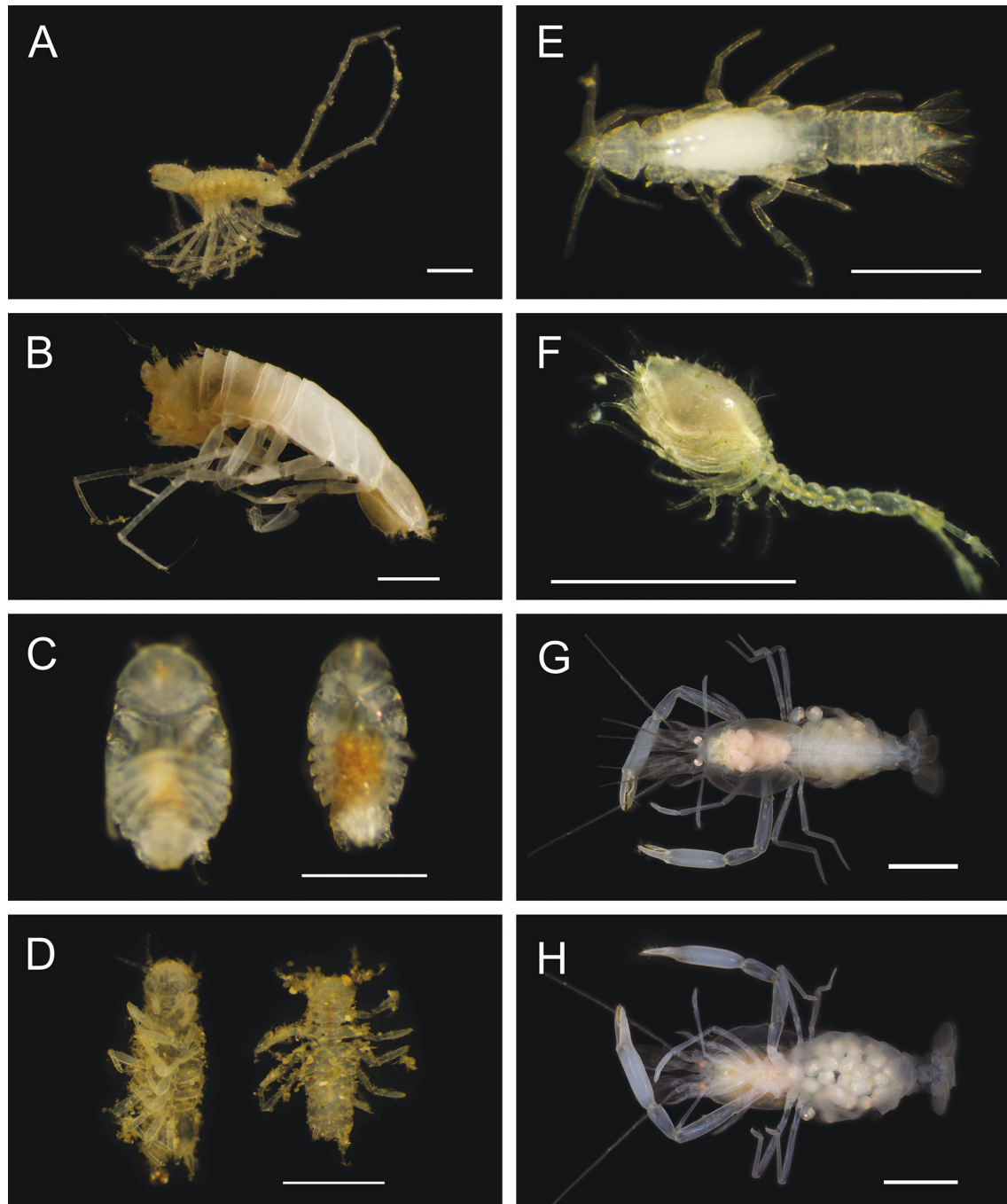


Figure 52. Arthropoda: Isopoda, Cumacea, and Stenopodidea, representative live images **A** *Munna* gen. inc. (C13872) **B** *Betamorpha* stet. (C13974) **C** *Eurycope* stet. (C13850) **D** *Tenupedunculus* stet. (C13823) **E** Gnathiidae stet. (C13801) **F** *Paraleucon* sp. SIO_BIC_C11152 (C11152) **G** *Spongiocoloides* cf. *galapagensis* (C12820, dorsal view) **H** *Spongiocoloides* cf. *galapagensis* (C12820, ventral view). Scale bars: 1 mm (**A–F**); 1 cm (**G, H**).

Arthropoda | ... | Decapoda | Pleocyemata | Caridea | Alpheoidea | Thoridae

***Lebbeus scrippsi* Wicksten & Méndez G., 1982**

Fig. 53A

Material examined. AD4924: C12822; AD4989: C13878, C13879, MZU-CR-3747-01; AD4990: C13880, C13881, C13882.

Localities. Parrita Seep (~ 1400–1410 m), Jacó Scar (~ 1768–1785 m).

Distribution. Originally described from northern Chile and southern Peru, 768–1164 m (Wicksten and Méndez 1982), and recorded as far north as the southeastern Gulf of California, 1188–1245 m (Hendrickx 1996, 2001).

New records. CRM specimens C13878, C13879, and MZUCR-3747-01 represent new maximum depth records for this species (1768 m as the most conservative value).

Arthropoda | ... | Decapoda | Pleocyemata | Caridea | Bresilioidea | Alvinocarididae

***Alvinocaris costaricensis* Martin, Wall, Shank, Cha, Seid & Rouse, 2018**

Fig. 53B

Reference. Martin et al. 2018**.

Additional material examined. S0213: MZUCR 3748-01 (voucher), C13898 (tissue).

Localities. Jacó Summit (742 m; this study), Mound 12 (~ 1000 m; type locality), Jacó Scar (~ 1800 m).

Distribution. Known only from the CRM seeps.

New records. Specimen MZUCR 3748-01 from Jacó Summit (742 m) represents a new minimum depth record for this species.

***Alvinocaris* sp. SIO_BIC_C11136**

Fig. 53C

Reference. Martin et al. 2018.

Material examined. AD4501: C11136; AD4503: C13307; AD4589: C11190; AD4974: C13782; AD4978: C13806; AD4984: C13861, C13862, C13863, C13893, MZUCR-3749-01.

Localities. Mound 12 (~ 967–999 m).

Remarks. An undescribed species, previously noted as morphologically and genetically distinct from *A. costaricensis* (Martin et al. 2018).

Arthropoda | ... | Decapoda | Pleocyemata | Caridea | Crangonoidea | Crangonidae

***Paracrangon areolata* Faxon, 1893**

Fig. 53D

Material examined. AD4924: C12751 ([PQ449342](#); 16S: [PQ304652](#); 18S: [PQ304647](#)).

Localities. Parrita Seep (1402 m).

Distribution. Southern Gulf of California and western Mexico (type locality: Islas Mariás, 1236–1244 m (Faxon 1893, 1895)) to central Chile, 580–1650 m (Hendrickx 1996; Báez and Soto 1997; Wicksten 2020).

Remarks. This specimen is an ovigerous female.

Arthropoda | ... | Decapoda | Pleocyemata | Caridea |
Nematocarcinoidea | Nematocarcinidae

***Nematocarcinus faxoni* Burukovsky, 2001**

Fig. 53E

Material examined. S0213: C13899.

Localities. Jacó Summit (763 m).

Distribution. Mexico to the Galápagos Islands and southwestern Atlantic, 660–2055 m (Hendrickx and Hernández-Payán 2018; Martínez-Guerrero and López-Pérez 2018).

Arthropoda | ... | Decapoda | Pleocyemata | Caridea | Pandalioidea |
Pandalidae

***Heterocarpus vicarius* Faxon, 1893**

Fig. 53F

Material examined. S0213: C13900; S0219: C13921.

Localities. Rio Bongo Scar (661 m), Jacó Summit (763 m).

Distribution. Originally described from several stations in the Gulf of Panama, 384–523 m (Faxon 1893, 1895) and recorded from the Gulf of California to Peru, 73–1454 m (Fischer et al. 1995). This species is commercially harvested in Costa Rica, typically at depths of 250–300 m (Fischer et al. 1995; Wehrtmann and Echeverría-Sáenz 2007).

Remarks. C13921 was associated with a naturally occurring wood fall.

***Plesionika trispinus* Squires & Barragan, 1976**

Fig. 53G

Material examined. AD4919: C12806.

Localities. Quepos Slide (~ 379–397 m).

Distribution. Originally described from 209–302 m off the Pacific coast of Colombia (Squires and Barragan 1976) and recorded from the southern Gulf of California to Peru, 96–500 m (Fischer et al. 1995), including the Galápagos Islands (Arnés-Urgellés et al. 2020) and Costa Rica (Wehrtmann and Echeverría-Sáenz 2007; Azofeifa-Solano and Cortés 2020).

Arthropoda | ... | Decapoda | Pleocyemata | Anomura | Chirostyloidea |
Chirostylidae

***Heteroptychus galapagos* Baba & Wicksten, 2019**

Figs 53H, 54A

Material examined. AD4923: C12816 ([PQ449348](#)), C12817 ([PQ449349](#)).

Localities. Parrita Seep (1037 m).

Distribution. Previously known only from East Darwin Seamount, Galápagos Islands, 1012 m (Baba and Wicksten 2019).

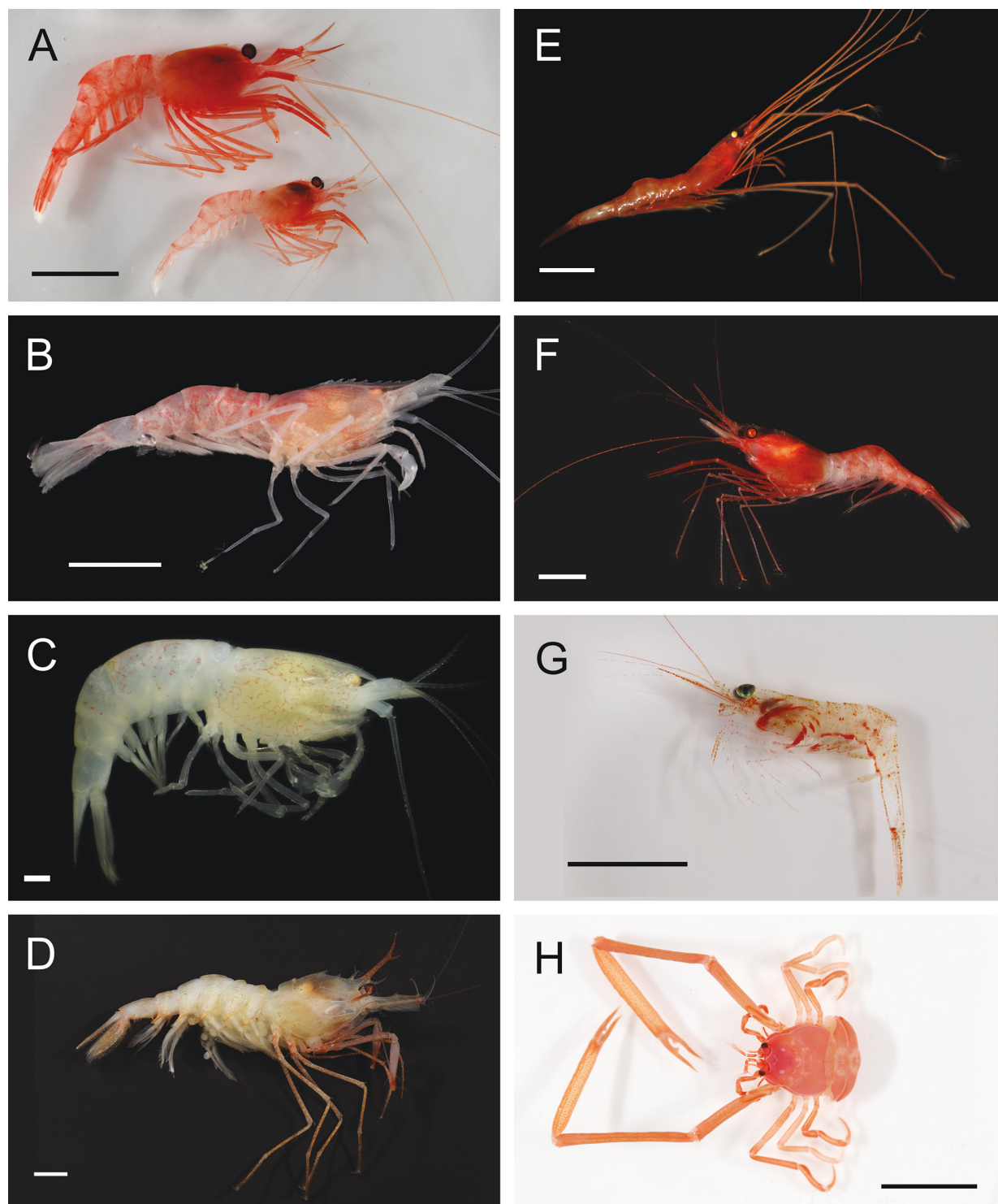


Figure 53. Arthropoda: Caridea and Anomura, representative live images **A** *Lebbeus scrippsi* (C12822) **B** *Alvinocaris co-staricensis* (MZUCR 3748-01) **C** *Alvinocaris* sp. SIO_BIC_C11136 (C11136) **D** *Paracrangon areolata* (C12751) **E** *Nematocarcinus faxoni* (C13899) **F** *Heterocarpus vicarius* (C13921) **G** *Plesionika trispinus* (C12806) **H** *Heteroptychus galapagos* (C12816, dorsal view). Scale bars: 1 cm (**A, B, D–H**); 1 mm (**C**).

New records. The CRM specimens represent new northern records of this species.

Remarks. Associated with isidid corals Co2952, Co2953, and/or MZUCR 3129. C12816 is an ovigerous female.

***Sternostylus defensus* (Benedict, 1902)**

Fig. 54B–D

Material examined. AD4923: C12804 ([PQ450385](#)).

Localities. Parrita Seep (~ 1035–1108 m).

Distribution. Originally described from the Galápagos Islands, 717 m (Baba and Haig 1990), and recently reported from 873–1210 m at seamounts in the northern Galápagos Marine Reserve (Arnés-Urgellés et al. 2020; Salinas-de-León et al. 2020).

New records. The CRM specimen represents a new northern record of this species.

Remarks. Several antipatharians and octocorals were collected during this dive, and the exact host was not recorded. At the Galápagos seamounts, *S. defensus* is likely associated with antipatharian corals and possibly with *Chrysogorgia* (Baba and Wicksten 2019).

Arthropoda | ... | Decapoda | Pleocyemata | Anomura | Chirostyloidea | Kiwaidae

***Kiwa puravida* Thurber, Jones & Schnabel, 2011**

Fig. 54E

Reference. Thurber et al. 2011**.

Localities. Mound 12 (~ 1000 m; type locality), Mound 11 (~ 1000–1040 m).

Distribution. Known only from the CRM seeps.

Remarks. An iconic megafaunal species of the CRM seeps (Fig. 2D), the “Costa Rican dancing yeti crab” cultivates ectosymbiotic bacteria on its chelipeds and harvests the bacteria as its main source of food; the animals are often found in aggregations at areas of active fluid seepage, where the slow waving of the chelipeds is hypothesized to shear off the nutrient-depleted boundary layer and thereby enhance the productivity of the ectosymbionts (Thurber et al. 2011).

Arthropoda | ... | Decapoda | Pleocyemata | Anomura | Galatheaidea | Munididae

***Grimothea monodon* (H. Milne Edwards, 1837)**

Fig. 54F

Material examined. AD4918: C12805; AD4921: C12808 ([PQ449345](#)); AD4979: C13809, C13810, C13811.

Localities. Quepos Slide (~ 333–408 m).

Distribution. Previously recorded from Costa Rica, 150–350 m, where it is abundant as bycatch in deepwater shrimp fisheries (Wehrtmann et al. 2010). Also known from central Mexico to southern Chile (type locality: unspecified site on the Chilean coast) (Milne Edwards 1837), to 523 m depth (Faxon 1895; Hendrickx and Harvey 1999; Baba et al. 2008).

Remarks. This species is commercially harvested off Chile and serves as prey for other species of commercial interest (Yapur-Pancorvo et al. 2023).

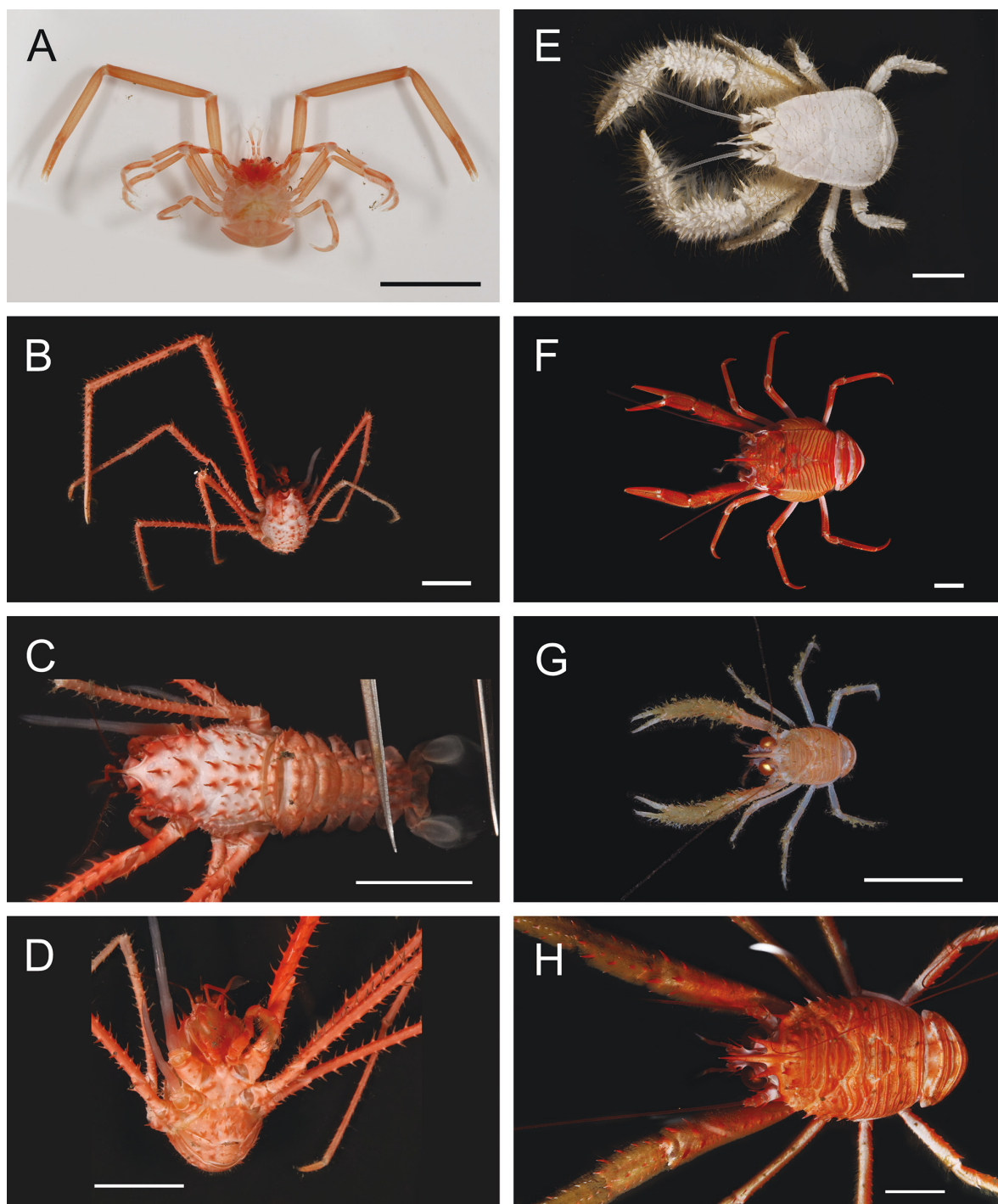


Figure 54. Arthropoda: Anomura, representative live images **A** *Heteroptychus galapagos* (C12817, ventral view) **B** *Sternostylus defensus* (C12804, dorsal view) **C** *Sternostylus defensus* (C12804, dorsal detail) **D** *Sternostylus defensus* (C12804, ventral view) **E** *Kiwa puravida* (MZUCR 3735-01) **F** *Grimothea monodon* (C13909) **G** *Typhlonida propinqua* (C11153, juvenile) **H** *Typhlonida propinqua* (C13915, adult). Scale bars: 1 cm.

***Typhlonida propinqua* (Faxon, 1893)**

Fig. 54G, H

Material examined. AD4510: C11153; S0213: C13901; S0215: C13915, MZUCR-3746-01; S0219: C13925.

Localities. Rio Bongo Scar (661 m), Jacó Summit (741–763 m), Mound 12 (991 m).

Distribution. Originally described from the Gulf of Panama (838–935 m) and Galápagos Islands (704 m) (Faxon 1895) and distributed south to Peru and central Chile, 700–1713 m (Wicksten 1989, 2020; Baba et al. 2008; Retamal et al. 2020).

Remarks. C11153, C13901, and C13925 are juveniles. C13925 was associated with a naturally occurring wood fall.

Arthropoda | ... | Decapoda | Pleocyemata | Anomura | Galattheoidea |
Munidopsidae

We thank Paula Rodríguez-Flores (Harvard University, U.S. National Museum of Natural History) for contributing to morphological identification of these specimens and generating many DNA sequences. See Rodríguez-Flores et al. (2023) for discussion of the phylogenetics and biogeography of eastern Pacific Munidopsidae.

***Munidopsis agassizii* Faxon, 1893**

Fig. 55A

Reference. Rodríguez-Flores et al. (2023) for phylogenetic analysis.

Material examined. S0219: C13922 ([ON886860](#)), C13923.

Localities. Rio Bongo Scar (661 m).

Distribution. Gulf of Panama (type locality) to northern Chile, 384–1000 m (Guzman and Sellanes 2015).

***Munidopsis alvisca* A.B. Williams, 1988**

Fig. 55B

Reference. Rodríguez-Flores et al. (2023) for phylogenetic analysis.

Material examined. AD4509: C11149; AD4513: C11158, C11159 ([PQ450374](#)); AD4914: C12783 ([ON886848](#)), C12790 ([ON886849](#)); AD4591: C11181; AD4914: C12791; AD4972: C13792; AD4989: C13876; AD4990: C13883; S0214: C13904.

Localities. Parrita Seep (1401 m; this study), Jacó Slope (1459 m; this study), Jacó Scar (~ 1632–1886 m).

Distribution. Originally described from vents in the Guaymas Basin (type locality), Gulf of California, 2008 m, and the Explorer and Juan de Fuca Ridges, 1545–1818 m (Williams 1988). Material identified as “most probably *Munidopsis alvisca*” has been reported from Pacific Costa Rica (locality and depth not specified) (Wehrtmann et al. 2010).

New records. Confirming the occurrence of *M. alvisca* in Costa Rica, our specimens represent new southern records, new seep records, and a new minimum depth record for this species (specimen C13883 from 1401 m).

Remarks. Several specimens are ovigerous females: C11149, C11158, C11181, and C13792. Specimen C13876 was the host of sacculinid parasite C13875.

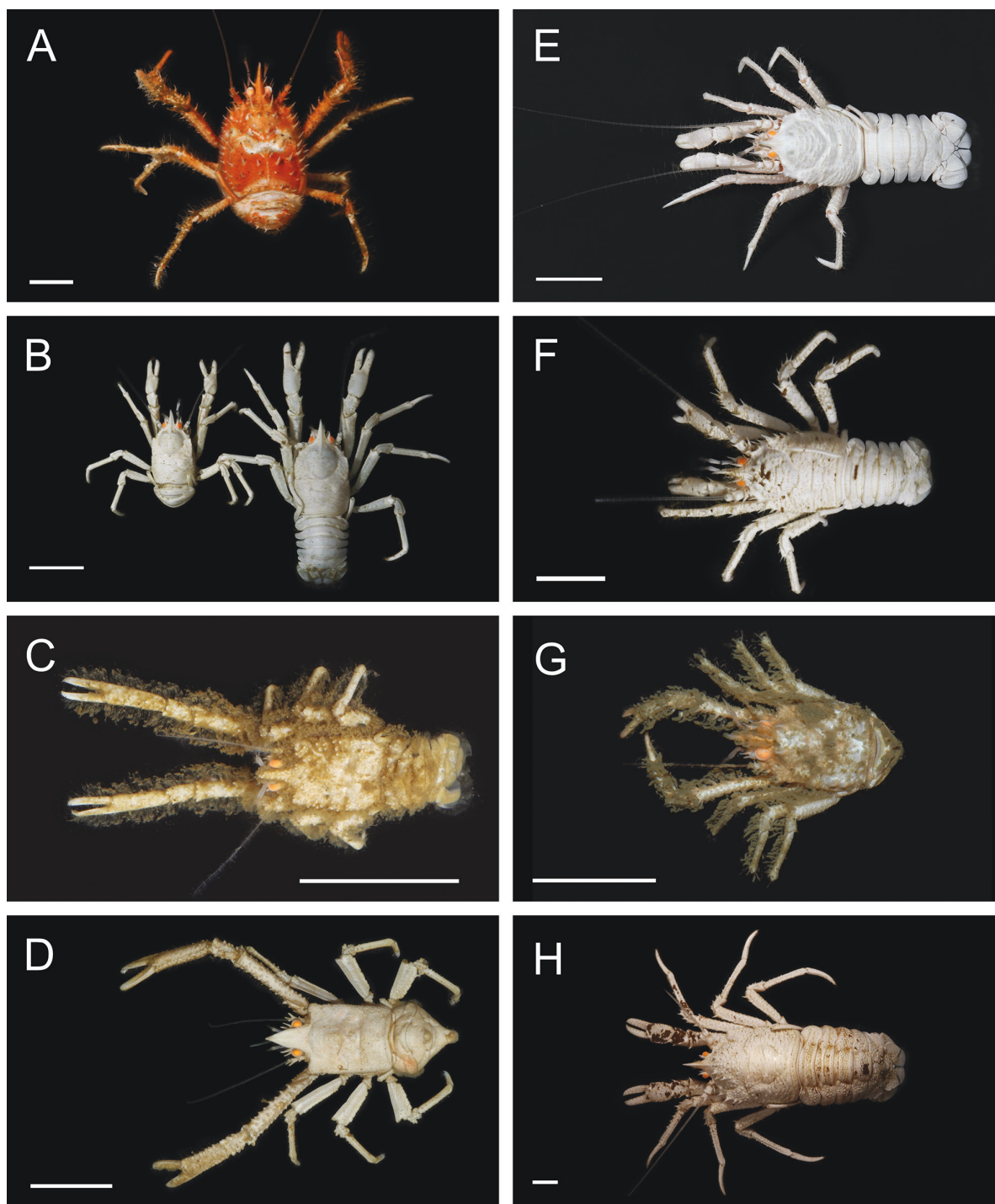


Figure 55. Arthropoda: *Munidopsis*, representative live images **A** *Munidopsis agassizii* (C13922) **B** *Munidopsis alvisca* (C13904) **C** *Munidopsis aspera* (C13912) **D** *Munidopsis carinipes* (C11150) **E** *Munidopsis* cf. *nitida* (C12818) **F** *Munidopsis* cf. *nitida* (C13903) **G** *Munidopsis* cf. *opalescens* (C11154) **H** *Munidopsis* cf. *producta* (MZUCR-3745-01). Scale bars: 1 cm.

***Munidopsis aspera* (Henderson, 1885)**

Fig. 55C

Reference. Rodríguez-Flores et al. (2023) for phylogenetic and taxonomic treatment.

Localities. The Thumb (1065 m).

Distribution. Originally described from the Straits of Magellan, 448 m (Henderson, 1885), and recorded along the eastern Pacific from southern California to northern Chile, 166–1398 m (Rodríguez-Flores et al. 2023), including records from Isla del Coco (Baba et al. 2008).

Remarks. Specimens C13912 and C13913 from dive S0217 were associated with the holothuroid E7313 (*Synallactes* cf. *chuni*) and heavily coated with bacteria and/or sediment.

***Munidopsis carinipes* Faxon, 1893**

Fig. 55D

Reference. Rodríguez-Flores et al. (2023) for phylogenetic and taxonomic treatment.

Localities. Mound 12 (~ 990–1011 m), Mound 11 (1010 m), Jacó Scar (1459 m).

Distribution. Known only from the CRM and western Panama, 915–1459 m (Rodríguez-Flores et al. 2023).

Remarks. Specimen C13869 from dive AD4988 was associated with a naturally occurring wood fall. Specimen C13916 from dive S0215 was associated with experimentally deployed cow or pig bones.

***Munidopsis* cf. *nitida* (A. Milne Edwards, 1880)**

Fig. 55E, F

Reference. Rodríguez-Flores et al. (2023) for phylogenetic analysis.

Material examined. AD4924: C12818 ([PQ450377](#)); S0214: C13903 ([ON886857](#); 16S: [ON858040](#); 28S: [ON858109](#)).

Localities. Parrita Seep (1402 m), Jacó Scar (1875 m).

Remarks. C13903 was associated with a naturally occurring wood fall. These specimens resemble *M. nitida*, but C13903 belongs to a potential species complex warranting further investigation (Rodríguez-Flores et al. 2023). Originally described from Guadeloupe Island, Caribbean Sea, 1407–1607 m, *M. nitida* has been reported from the western Atlantic, Indian Ocean, western Pacific, and eastern Pacific over a depth range of 592–3680 m (Baba et al. 2008). In the eastern Pacific, *M. nitida* has been reported from the Gulf of Panama and Isla del Coco (Wehrtmann et al. 2010; Azofeifa-Solano and Cortés 2020; Wicksten 2020).

***Munidopsis* cf. *opalescens* Benedict, 1902**

Fig. 55G

Reference. Rodríguez-Flores et al. (2023) for phylogenetic analysis (reported as *M. opalescens*).

Material examined. AD4510: C11154 ([ON886842](#)).

Localities. Jacó Summit (741 m).

Remarks. *M. opalescens* was originally described from the Magellan region of Chile and is known only from Chilean waters, 636–922 m (Wicksten 1989, 2020; Baba et al. 2008; Guzman and Sellanes 2015), with a northernmost record at 32°S (Guzman and Sellanes 2015). Although the CRM specimen is morphologically

consistent with *M. opalescens*, the COI sequence was only 98.68% identical to a reference sequence of *M. opalescens* from Chile ([JN166759.1](#)) (Ahyong et al. 2011), exceeding the typically low (<1%) intraspecific variation in Pacific *Munidopsis* species (Jones and Macpherson 2007; Rodríguez-Flores et al. 2023). If confirmed as *M. opalescens* by further comparison to type material, the CRM specimen would represent the first record of *M. opalescens* in the northern hemisphere and a northward range extension of ca 5000 km.

***Munidopsis cf. producta* Baba, 2005**

Fig. 55H

Material examined. S0220: MZUCR-3745-01 (voucher), C13926 (tissue).

Localities. Subduction Plume (3399 m).

Remarks. *M. producta* was originally described from Costa Rica (ca 400 km offshore, 3680 m) and occurs from the Pescadero Basin hydrothermal vents in the Gulf of California (Rodríguez-Flores et al. 2023) to the Gulf of Panama, 3260–3680 m (Baba 2005; Wicksten 2020). The CRM specimen, an ovigerous female, appears morphologically consistent with *M. producta* and was collected ~ 500 km from the type locality at a similar depth. DNA sequences were not obtained due to bacterial contamination and more detailed morphological comparison is required, so we conservatively designate this specimen as *M. cf. producta*.

***Munidopsis cf. trifida* Henderson, 1885**

Fig. 56A, B

Reference. Rodríguez-Flores et al. (2023) for phylogenetic analysis (reported as *M. trifida*).

Material examined. AD4507: C11145 ([ON886840](#)); AD4586: C14391; AD4923: C12813 ([ON886850](#); 16S: [ON858033](#); 28S: [ON858102](#)); AD4985: C13829 (tissue); S0217: MZUCR-3733-01 (voucher), C13918 (tissue).

Localities. Mound 12 (~ 982–998 m), The Thumb (1071 m), Parrita Seep (1091 m), Parrita Scar (1660 m).

Remarks. *M. trifida* was originally described from the Straits of Magellan, Chile, 732 m (Henderson 1885), and has been reported from the western Pacific and Indian Oceans, 280–1100 m (Baba 2005; Baba et al. 2008). The COI sequence of C11145 was identical to those of specimens from Chile that were previously identified as *M. trifida* ([JN166765.1](#), [JN166764.1](#)) (Ahyong et al. 2011). Cryptic diversity within this putatively widespread species is currently under investigation, and the CRM specimens may represent at least one undescribed species. Any CRM specimens confirmed as *M. trifida sensu stricto* would represent new northern records in the eastern Pacific and new depth records for this species.

***Munidopsis cortesi* Rodríguez-Flores, Seid, Rouse & Giribet, 2023**

Fig. 56C

Reference. Rodríguez-Flores et al. 2023**.

Localities. Jacó Summit (742 m; type locality).

Distribution. Known only from the CRM seeps.

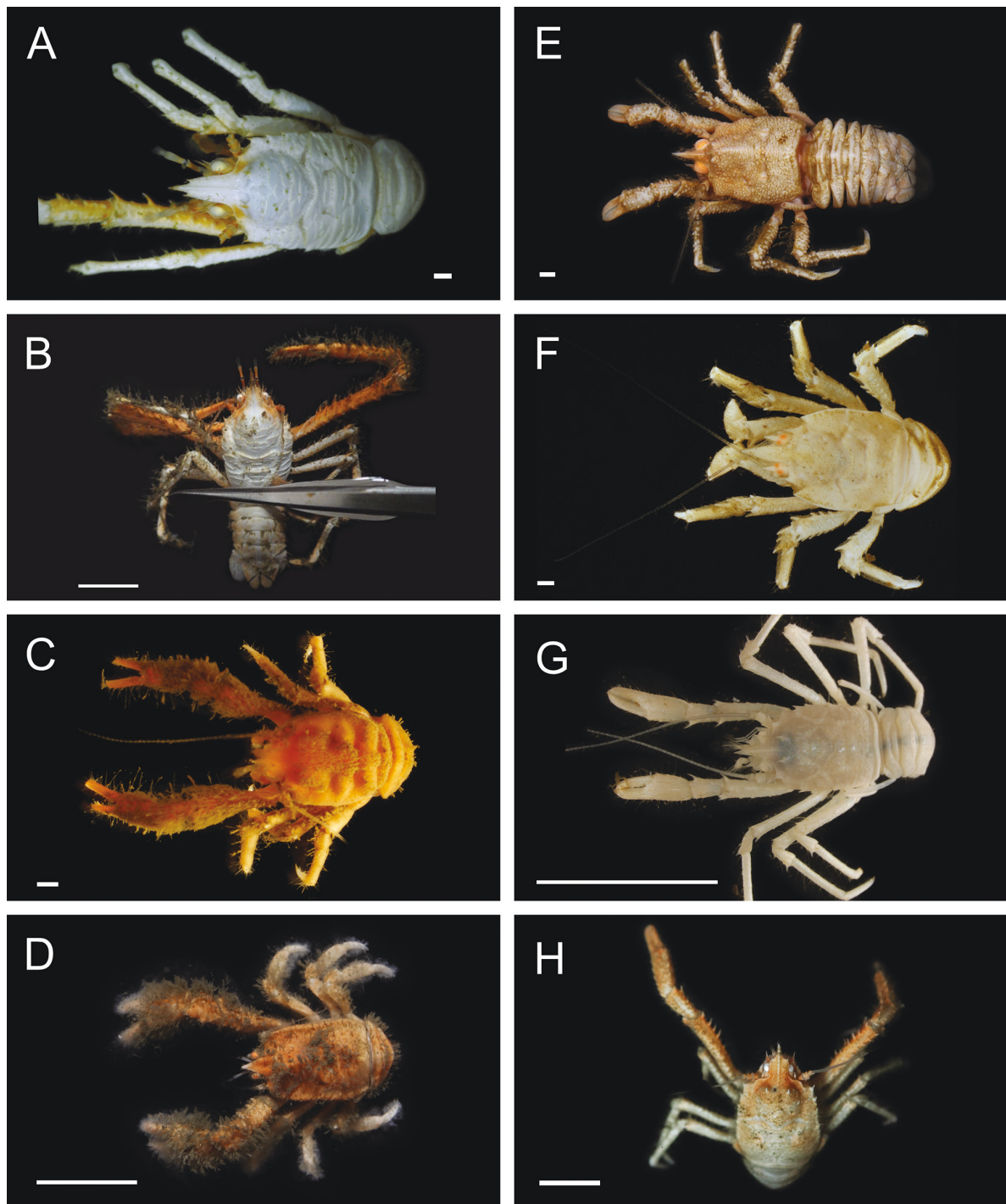


Figure 56. Arthropoda: *Munidopsis*, representative images. Live specimens are depicted unless otherwise specified **A** *Munidopsis* cf. *trifida* (C11145) **B** *Munidopsis* cf. *trifida* (C12813) **C** *Munidopsis* *cortesi* (C13902) **D** *Munidopsis* *girguisi* (C13897) **E** *Munidopsis* *granosicorium* (C12819) **F** *Munidopsis* *hendersoniana* (C12810) **G** *Munidopsis* *similis* (C13964, preserved specimen, credit: Paula Rodríguez-Flores) **H** *Munidopsis* sp. SIO_BIC_C11177 (C13830). Scale bars: 1 mm (A, C, E, F); 1 cm (B, D, G, H).

***Munidopsis girguisi* Rodríguez-Flores, Seid, Rouse & Giribet, 2023**

Fig. 56D

Reference. Rodríguez-Flores et al. 2023**.

Localities. Jacó Summit (~ 730–820 m).

Distribution. Eastern Pacific seeps from southern California (type locality) to the CRM, 381–845 m (Rodríguez-Flores et al. 2023).

***Munidopsis granosicorium* A.B. Williams & Baba, 1989**

Fig. 56E

Reference. Rodríguez-Flores et al. (2023) for phylogenetic and taxonomic treatment.

Localities. Parrita Seep (1402 m).

Distribution. Cascadia Basin off the Strait of Juan de Fuca (type locality) to the CRM, 1402–2020 m (Rodríguez-Flores et al. 2023).

***Munidopsis hendersoniana* Faxon, 1893**

Fig. 56F

Reference. Rodríguez-Flores et al. (2023) for phylogenetic analysis.

Material examined. AD4503: C11142 ([ON886839](#)); AD4922: C12803, C12809 ([ON886884](#)), C12810, C12812; AD4974: C13783; AD4988: C13868.

Localities. Mound 12 (~ 966–1005 m), Mound 11 (1010 m).

Distribution. Southwestern Mexico to the Gulf of Panama (type locality), 1101–1869 m (Faxon 1895; Baba 2005; Wicksten 2020).

Remarks. Some specimens were associated with naturally occurring wood falls (C11142, C12810, and C13868) or cnidarians (C12799 with hormathiid anemone Co2933; C12803 with hormathiid anemone Co2875; C12809 with anthoptilid sea pen Co2937). C12809 includes an ovigerous female.

***Munidopsis similis* Smith, 1885**

Fig. 56G

Reference. Rodríguez-Flores et al. (2023) for phylogenetic and taxonomic treatment.

Localities. Mound Jaguar (1895 m).

Distribution. Originally described from the North Atlantic off the eastern United States, 1939 m (Smith 1885), with apparent genetic connectivity from the northwest Atlantic Ocean to the northeast Pacific and Bering Sea, 1390–3314 m (Rodríguez-Flores et al. 2023).

***Munidopsis* sp. SIO_BIC_C11177**

Fig. 56H

Reference. Rodríguez-Flores et al. (2023) for phylogenetic analysis.

Material examined. AD4590: C11173, C11175, C11177 ([PQ450376](#)), C11179 ([PQ450375](#)); AD4985: C13830 ([ON886853](#)).

Localities. Mound 12 (~ 1000 m), Jacó Scar (1800 m).

Remarks. An undescribed species.

Arthropoda | ... | Decapoda | Pleocyemata | Anomura | Lithodoidea |
Lithodidae

***Glyptolithodes* aff. *crisatipes* (Faxon, 1893)**

Fig. 57A, B

Material examined. AD4916: C12793 (tissue).

Localities. Jacó Scar (1742 m).

Remarks. Resembles but is distinct from *Glyptolithodes crisatipes* (Faxon, 1893), which was described from western Panama, 589 m (Faxon 1895).

***Lithodes* cf. *couesi* Benedict, 1895**

Fig. 57C

Material examined. AD4985: C14058 ([PQ449358](#)).

Localities. Mound 12 (995 m depth, ~ 5 m above the seafloor).

Remarks. *L. couesi* was originally described from Unalaska, Aleutian Islands, at 730 m (Benedict, 1895), and has been reported to range from northern Japan to Baja California, with an overall depth range of 258–1400 m (Wicksten, 1989; Hendrickx et al. 2014; Wicksten, 2020). A recent molecular study reported *L. couesi* from the Burdwood Bank, Scotia Arc, southwestern Atlantic Ocean, 605 m, showing only minimal COI haplotype variation compared to individuals from Alaska and British Columbia and indicating a pole-to-pole distribution of this species (Pérez-Barros et al. 2020). Pérez-Barros et al. (2020) hypothesize that *L. couesi* colonized the Burdwood Bank via deep-sea larval circulation, and that with increased sampling effort *L. couesi* is likely to be reported from intermediate locations between Baja California and the Scotia Arc.

Consistent with this hypothesis, the COI sequence of the CRM lithodid larva C14058 is 100% identical to that of *L. couesi* from British Columbia ([DQ882085.1](#)) and only 2–3 bases different from additional *L. couesi* sequences ([DQ882086.1](#) from British Columbia, 99.53% identity; [MN047311.1](#) from Burdwood Bank, 99.83% identity) (Fig. 57D). The C14058 sequence is also only 4 bases (<1%) divergent from that of the CRM *Lithodes panamensis* specimen C13787 (below), which represents the first available reference sequence for *L. panamensis*. Evidently, COI does not meaningfully differentiate between these species, and further investigation or revision is warranted.

***Lithodes panamensis* Faxon, 1893**

Fig. 57E, F

Material examined. AD4975: C13787 ([PQ449352](#)).

Localities. Mound 12 (1000 m). Previously reported off Puntarenas, Costa Rica, 700–1400 m, although without specific mention of a seep habitat (MacPherson and Wehrtmann 2010).

Distribution. Originally described from 838 m in the Gulf of Panama (Faxon 1893, 1895) and reported to the Galápagos seamounts, Peru, and northern Chile, 620–850 m (Wicksten 1989; Arguelles et al. 2020; Arnés-Urgellés et al. 2020; Retamal et al. 2020).

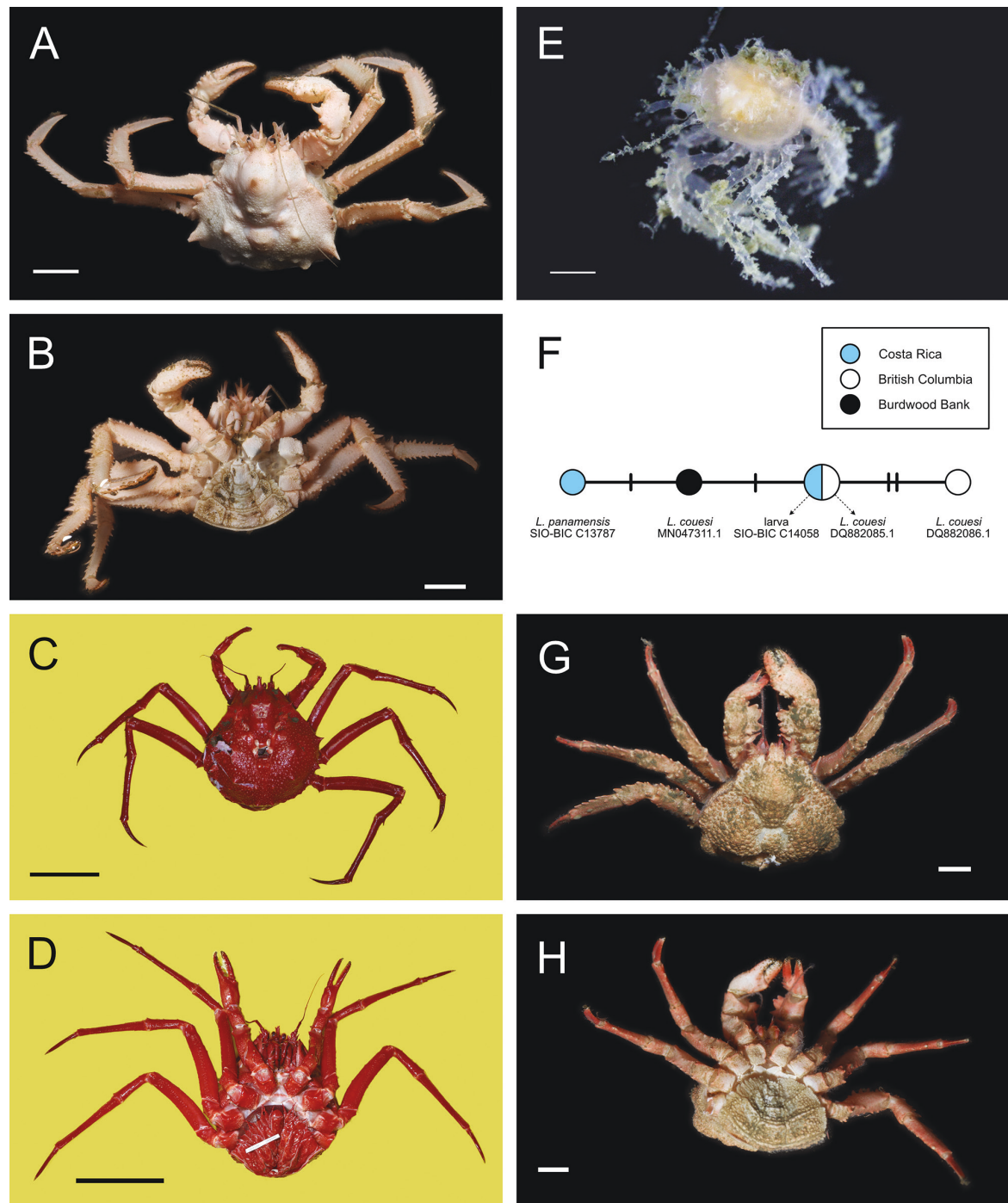


Figure 57. Arthropoda: Lithodidae, representative live images **A** *Glyptolithodes* aff. *cristatipes* (C12793, dorsal view) **B** *Glyptolithodes* aff. *cristatipes* (C12793, ventral view) **C** *Lithodes panamensis* (C13787, dorsal view) **D** *Lithodes panamensis* (C13787, ventral view) **E** *Lithodes* cf. *couesi* (C14058) **F** Haplotype network of *L. panamensis* and *L. couesi* COI sequences **G** *Paralomis* aff. *diomedeeae* (C13788, dorsal view) **H** *Paralomis* aff. *diomedeeae* (C13788, ventral view). Scale bars: 1 cm (**A**, **B**, **G**, **H**); 10 cm (**C**, **D**); 1 mm (**E**).

Remarks. Host of the rhizocephalan parasite C13776. In regions of Peru where *L. panamensis* co-occurs with *L. wiracocha* Haig, 1974 and *Paralomis longipes* Faxon, 1893, rhizocephalans infest *P. longipes* with a prevalence of 7%, whereas the other two species have not been observed with parasites, suggesting a possible host preference (Arguelles et al. 2020). Our specimen confirms that *L. panamensis* is susceptible to rhizocephalan parasitization, at least at the CRM.

***Paralomis* aff. *diomedae* (Faxon, 1893)**

Fig. 57G, H

Reference. Niemann et al. (2013) for occurrences at Mound 12.

Material examined. AD4502: C11137 ([PQ450391](#)); AD4586: C14950; AD4917: C12795; AD4975: C13788 ([PQ450383](#)); S0217: C13914.

Localities. Mound 12 (~ 987–1000 m), The Thumb (1072 m; this study).

Remarks. The voucher specimens above match the description and locality of a *Paralomis* morphospecies that has been observed grazing on microbial mats at Mound 12 (Niemann et al. 2013). This likely undescribed species is morphologically similar to *P. diomedae* but differs “by the granules on the dorsal carapace surface and the armature of the chelipeds and walking legs” (Niemann et al. 2013). Our specimens share these notable differences from *P. diomedae*, which is known from California to the Gulf of Panama (type locality) to Peru, 680–935 m, and has been previously reported from Costa Rica, 770–825 m (MacPherson and Wehrtmann 2010; Hendrickx and López 2020).

Arthropoda | ... | Decapoda | Pleocyemata | Anomura | Paguroidea |
Parapaguridae

We thank Rafael Lemaitre (U.S. National Museum of Natural History) for identification of these specimens.

***Oncopagurus haigae* (de Saint Laurent, 1972)**

Fig. 58A

Material examined. S0216: C13911.

Localities. Quepos Slide (312 m).

Distribution. Eastern Pacific from southern California (type locality) to southern Chile (44°S), 55–993 m (Lemaitre 2014).

Remarks. This specimen provides the first record of live color for this species.

***Parapagurus foraminosus* Lemaitre, 1999**

Fig. 58B, C

Material examined. AD4590: C11210; AD4591: C13308; AD4916: C12787.

Localities. Jacó Scar (~ 1800 m).

Distribution. Northern Mexico to Ecuador, including Isla del Coco and the Galápagos Islands (type locality), 915–2807 m (Hendrickx and Ayon-Parente 2009).

Remarks. Consistent with previous reports of *P. foraminosus* inhabiting gastropod shells completely covered by actinians (Hendrickx and Ayon-Parente 2009), these specimens were associated with *Paracalliactis* anemones (C11210 with Co2300, C13308 with Co2320, and C12787 with Co2924). C11210 is an ovigerous female.

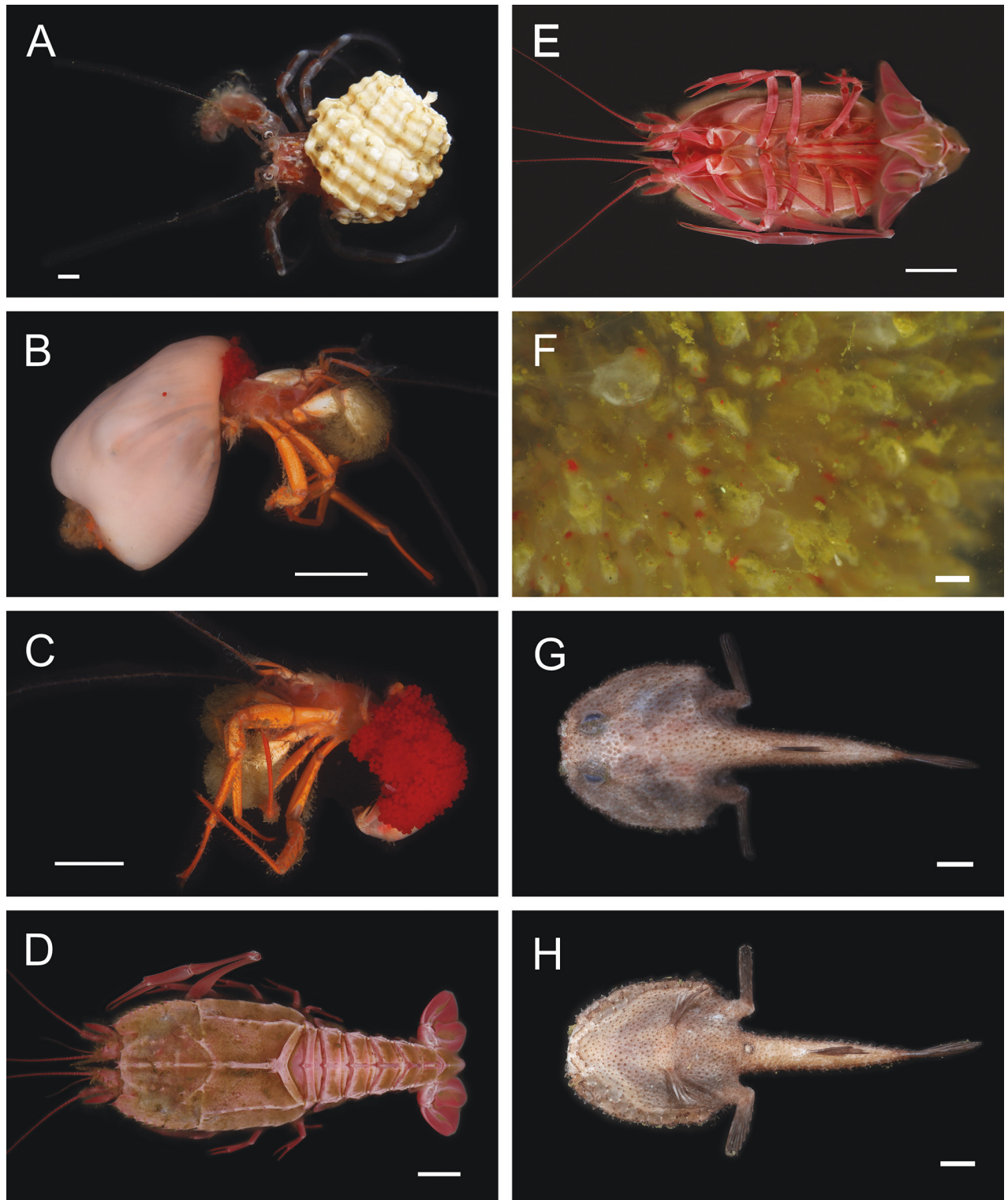


Figure 58. Arthropoda: Decapoda and Chordata, representative live images **A** *Oncopagurus haigae* (C13911) **B** *Parapagurus foraminosus* (C11210, dorsal view) **C** *Parapagurus foraminosus* (C11210, ventral view) **D** *Stereomastis pacifica* (C13962, dorsal view) **E** *Stereomastis pacifica* (C13962, ventral view) **F** Pyrosomatidae stet. (B11641) **G** *Dibranchius nudivomer* (UCR-FL-065, dorsal view) **H** *Dibranchius nudivomer* (UCR-FL-065, ventral view). Scale bars: 1 mm (**A**, **F**); 1 cm (**B**–**E**, **G**, **H**).

Arthropoda | ... | Decapoda | Pleocyemata | Polychelida | Polychelidae

***Stereomastis pacifica* (Faxon, 1893)**

Fig. 58D, E

Material examined. S0230: C13962 ([PQ449357](#)).

Localities. Mound Jaguar (1896 m).

Distribution. Originally described from the Gulf of Panama (type locality) and several stations off central and southern Mexico, 935–2323 m (Faxon 1893, 1895; Galil 2000). Recorded from northern California to central Chile, 600–3382 m, including previous records from Costa Rica (Galil 2000; Wicksten 2020). The large-bodied larvae occur in midwater to at least 3692 m (Wicksten 1980).

Chordata

Chordata | Tunicata | Thaliacea | Pyrosomatida

Pyrosomatidae stet.

Fig. 58F

Material examined. AD4512: BI1641.

Localities. Quepos Slide (400 m).

Chordata | Vertebrata | Actinopterygii | Lophiiformes | Ogcocephalidae

***Dibranchius nudivomer* (Garman, 1899)**

Fig. 58G, H

Material examined. S0217: UCR-FL-065 (voucher), ex BI1363 (tissue).

Localities. The Thumb (1065 m).

Distribution. Originally described from Pacific Colombia, 1271 m (Bradbury 2003), *D. nudivomer* is known from southern Baja California to Peru, 605–1457 m, and has been collected from Pacific Panama at depths of 1101–1314 m (Robertson et al. 2017).

Remarks. We thank Ben Fable (Scripps Institution of Oceanography Marine Vertebrate Collection) for the identification of this specimen.

Chordata | Vertebrata | Actinopterygii | Perciformes | Zoarcidae

***Pyrolycus jaco* Fable, Seid, Bronson & Møller, 2023**

Fig. 59A

Reference. Fable et al. 2023**.

Additional material examined. AD4916: BI1159 (tissue).

Localities. Jacó Scar (~ 1604–1854 m; type locality).

Distribution. Known only from the CRM seeps.

Remarks. Associated with *Lamellibrachia barhami* and *Escarpia spicata* tubeworm bushes (Fable et al. 2023).

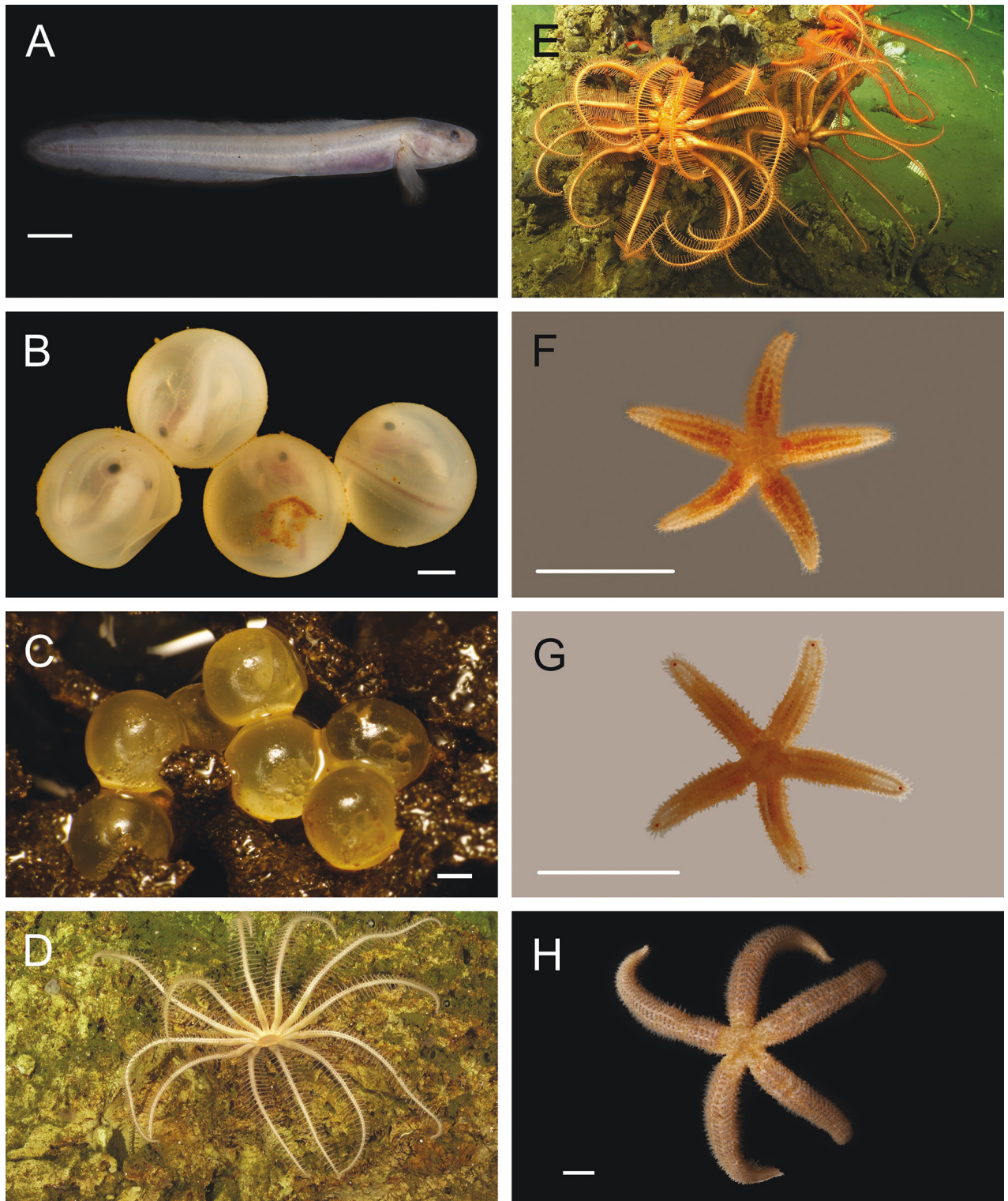


Figure 59. Chordata and Echinodermata: Asterozoidea, representative live images **A** *Pyrolycus jaco* (BI1159) **B** *Acantholiparis* stet. (BI1371, eggs) **C** *Paraliparis* stet. (BI1369, eggs) **D** *Freyella* stet. (E7365, *in situ*). Credit: ROV SuBastian/Schmidt Ocean Institute **E** *Freyella* stet. (E7367, *in situ*). Credit: ROV SuBastian/Schmidt Ocean Institute **F** *Ampheraster* gen. inc. (E7325, aboral view) **G** *Ampheraster* gen. inc. (E7325, oral view) **H** *Pedicellaster* gen. inc. (E7320, aboral view). Scale bars: 1 cm (**A**, **F–H**); 1 mm (**B**, **C**).

Chordata | Vertebrata | Actinopterygii | Scorpaeniformes | Liparidae

***Acantholiparis* stet.**

Fig. 59B

Reference. Levin and Rouse 2020.

Localities. Mound Jaguar (1903 m).

Remarks. This cluster of eggs, with visible embryos inside, was associated with a xenophyophore, *Shinkaya* stet., representing one of the first two recorded uses of xenophyophores as nursery habitat by fishes (Levin and Rouse 2020).

***Paraliparis* stet.**

Fig. 59C

Reference. Levin and Rouse 2020.

Localities. Jacó Scar (1866 m).

Remarks. This cluster of eggs was associated with a xenophyophore, *Reticulammina* stet., representing one of the first two recorded uses of xenophyophores as nursery habitat by fishes (Levin and Rouse 2020).

Echinodermata

We list the major clades according to the phylogenetic relationships in Reich et al. (2015).

Echinodermata | Asteroidea

We list entries following the phylogenetic relationships in Linchangco et al. (2017).

Echinodermata | Asteroidea | Brisingida | Freyellidae

***Freyella* stet.**

Fig. 59D, E

Material examined. S0230: E7365 ([PQ449392](#); arm fragment only), E7366 ([PQ449393](#); juvenile), E7367 ([PQ449394](#); arm fragment only).

Localities. Mound Jaguar (1895–1908 m).

Remarks. Morphologically, these specimens are most likely *Freyella*, although *Astrocles* and *Freyellaster* are possible and *Freyella* is not monophyletic (Zhang et al. 2024). The COI sequences were 98.97–99.13% identical to specimens of *Freyella* cf. *fragilissima* (Sladen, 1889) from Terre Adélie, Southern Ocean, 1956–2154 m ([OR600610.1](#), [OR600611.1](#)), and the Manus Basin, tropical western Pacific, 1708 m ([OR600608.1](#)).

Echinodermata | Asteroidea | Forcipulatida | Pedicellasteridae

***Ampheraster* gen. inc.**

Fig. 59F, G

Material examined. S0219: E7325 ([PQ449389](#); tissue).

Localities. Rio Bongo Scar (631 m).

Remarks. Juvenile pedicellasterid, likely *Ampheraster* based on spines along the marginals. The closest COI BLASTN results on GenBank were relatively distant, e.g., *Pedicellaster magister* Fisher, 1923 from British Columbia ([HM473941.1](#), 87.80% identity).

***Pedicellaster* gen. inc.**

Figs 59H, 60A

Material examined. S0219: E7320 ([PQ449388](#); tissue).

Localities. Rio Bongo Scar (609 m).

Remarks. The closest COI BLASTN result on GenBank was a specimen of *Ampheraster marianus* (Ludwig, 1905) from British Columbia ([HM542909.1](#), 94.28% identity).

Echinodermata | Asteroidea | Velatida | Xyloplacidae

***Xyloplax princealberti* Payne et al., 2023**

Fig. 60B

Reference. Payne et al. 2023**.

Localities. Jacó Scar (1845 m).

Distribution. Juan de Fuca Ridge (type locality) to the CRM seeps, on experimental wood and bone deployments and on a *Ridgea piscesae* Jones, 1985 tubeworm bush, 1845–2421 m (Payne et al. 2023).

***Xyloplax* sp. SIO_BIC_E7369**

Fig. 60C, D

Material examined. AD4972: E7223; AD4976: E7240, E7241; S0230: E7369.

Localities. Jacó Scar (1845–1887 m), Mound Jaguar (2000 m).

Remarks. An undescribed species, associated with naturally occurring and experimentally deployed wood, to be described in a separate work.

Echinodermata | Asteroidea | Paxillosida | Astropectinidae

***Tethyaster canaliculatus* (A.H. Clark, 1916) sp. inc.**

Fig. 60E, F

Material examined. AD4913: E7036.

Localities. Jacó Scar (1878 m).

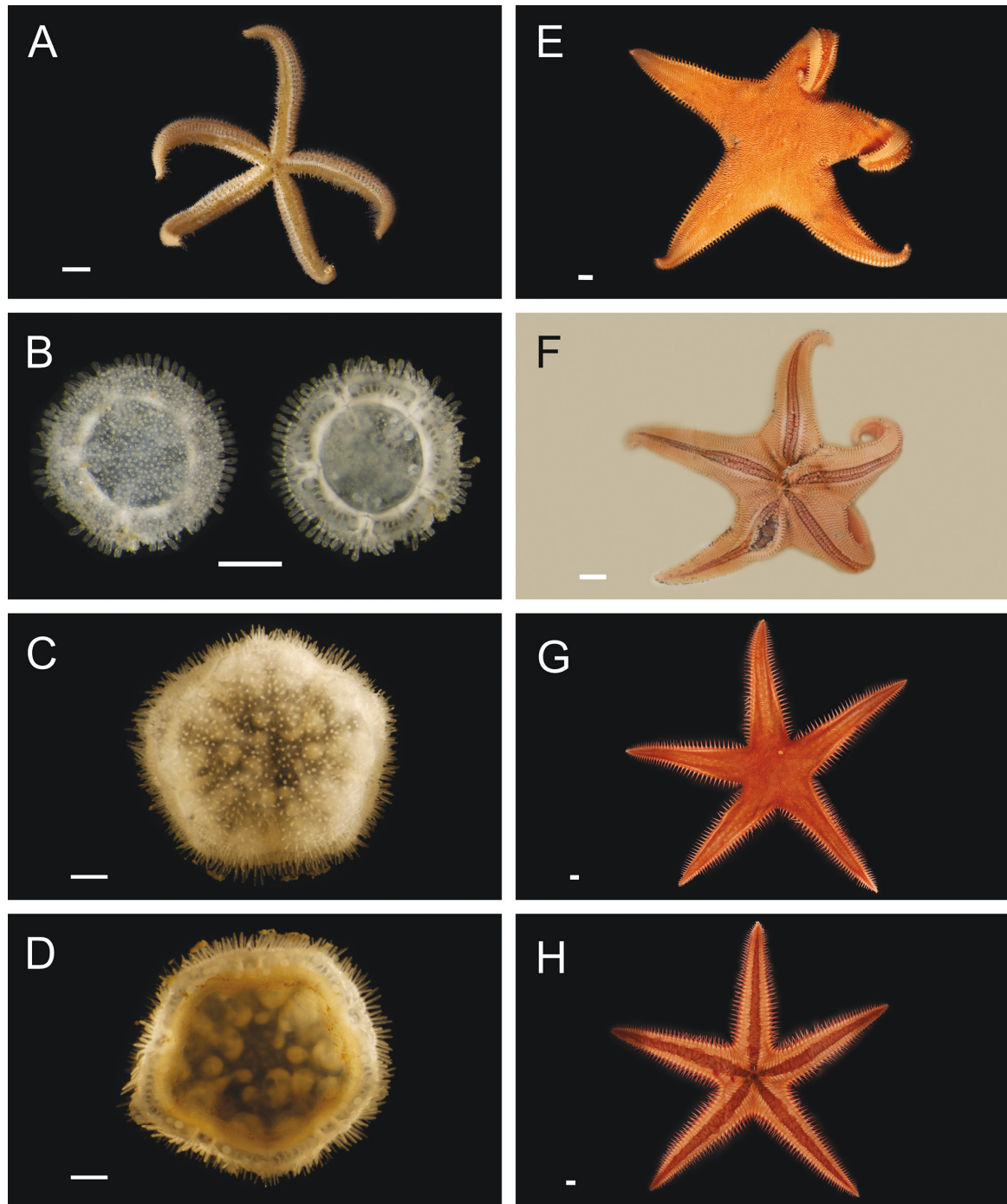


Figure 60. Echinodermata: Asterozoidea, representative live images **A** *Pedicellaster* gen. inc. (E7320, oral view) **B** *Xyloplax princealberti* (E7224, aboral and oral views) **C** *Xyloplax* sp. SIO_BIC_E7369 (E7369, aboral view) **D** *Xyloplax* sp. SIO_BIC_E7369 (E7369, oral view) **E** *Tethyaster canaliculatus* sp. inc. (E7036, aboral view) **F** *Tethyaster canaliculatus* sp. inc. (E7036, oral view) **G** *Thrissacanthias penicillatus* (E7246, aboral view) **H** *Thrissacanthias penicillatus* (E7246, oral view). Scale bars: 1 cm (**A**, **E–H**); 1 mm (**B–D**).

Remarks. Further morphological comparison is needed. *T. canaliculatus* was originally described from the Gulf of California at 73 m (Clark 1916a) and has been reported from Mexico to Peru at depths of 23–300 m (Alvarado and Solís-Marín 2013). If confirmed as *T. canaliculatus*, the CRM specimen would represent a new depth record for the species.

***Thrissacanthias penicillatus* (Fisher, 1905)**

Fig. 60G, H

Material examined. AD4975: E7246 (16S: [PQ304658](#)).

Localities. Mound 12 (1000 m).

Distribution. Originally described from the Coronados Islands off northern Mexico, 969–1167 m (Fisher 1905), *T. penicillatus* is known from Washington to the Gulf of California at depths of 55–1503 m (Fisher 1911; Hendrickx et al. 2011; Alvarado and Solís-Marín 2013).

New records. The CRM specimen represents a new southern record for the species. Its distribution may extend farther south, pending an unpublished report of *T. penicillatus* from Peru (“Morales-Montecinos (in preparation)” in Hooker et al. (2014)) and further identification of a specimen of *Thrissacanthias* sp. (MZUCR) from Pacific Costa Rica (Alvarado et al. 2017).

Remarks. This specimen was the host of at least six *Macellicephala* scale-worms (A9775) associated with the ambulacral groove. A COI sequence could not be obtained, but the closest 16S BLASTN results on GenBank showed 100.00% identity to reference sequences of *T. penicillatus* from California: [EU722968.1](#) (CASIZ 115075; off Mendocino County, 969 m) and [FJ177631.1](#) (SIO-BIC E3857, San Diego Trough, 1215 m).

Echinodermata | Asteroidea | Valvatida | Goniasteridae

***Ceramaster* gen. inc.**

Fig. 61A, B

Material examined. AD4917: E7048.

Localities. Mound 12 (997 m).

Remarks. Most likely *Ceramaster*, but internal dissection is required to exclude *Mediaster*.

***Evoplosoma claguei* Mah, Nizinski & Lundsten, 2010 sp. inc.**

Fig. 61C, D

Material examined. AD4923: E7081.

Localities. Parrita Seep (~ 1037–1108 m).

Distribution. *Evoplosoma claguei* is known from the Juan de Fuca Ridge, Rodriguez Seamount (type locality) off southern California, and Islas Tres Marías off central Mexico; 730–2406 m (Mah et al. 2010).

Remarks. If confirmed as *E. claguei*, the CRM specimen would represent a new southern record for the species.

***Nymphaster diomedeeae* Ludwig, 1905**

Fig. 61E, F

Reference. Mah (2016) for morphological examination of specimen E4386.

Material examined. AD4504: E4386 ([PQ449001](#)); AD4907: E7028; AD4917: E7051; AD4923: E7067, E7069; AD4975: E7235.

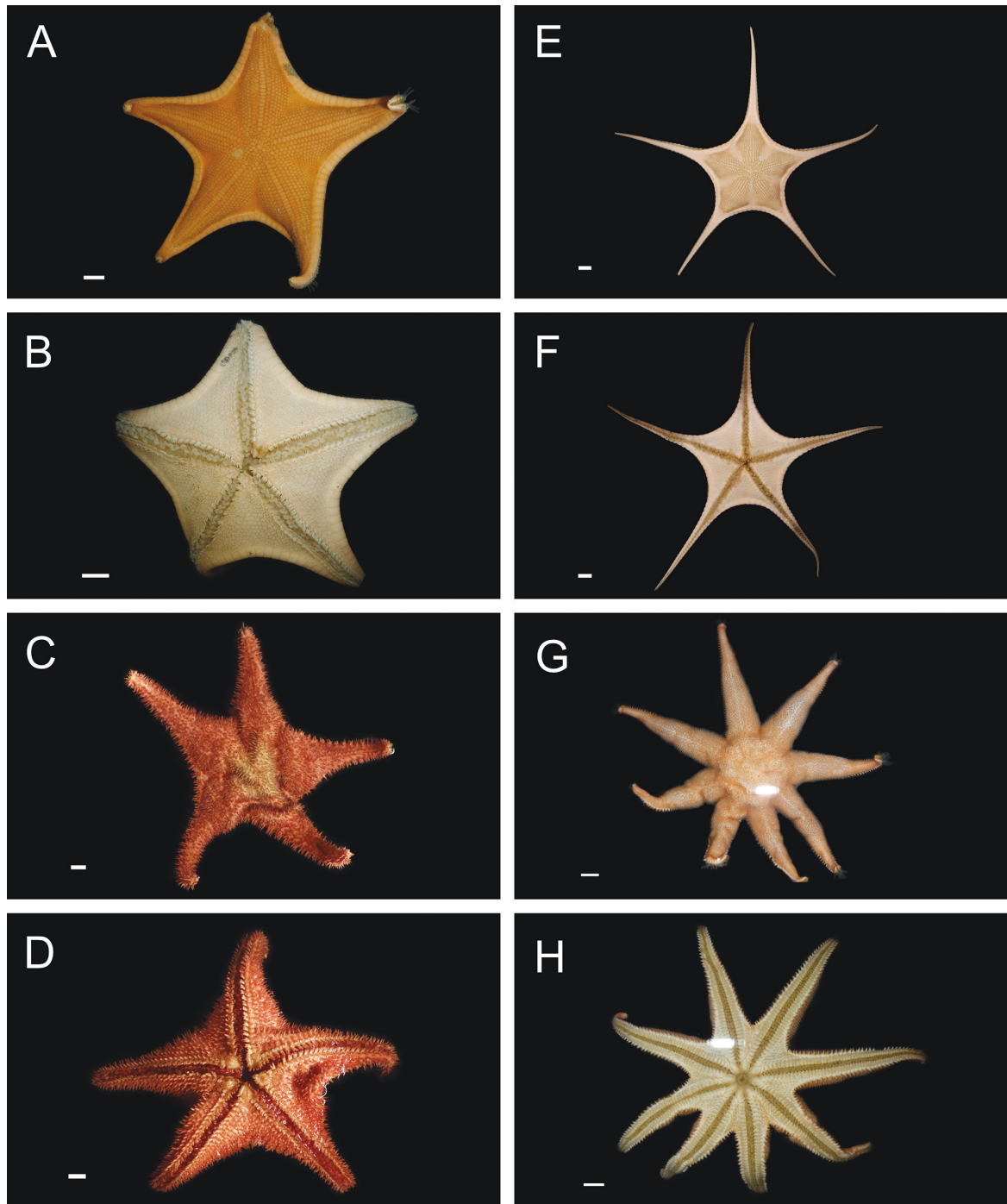


Figure 61. Echinodermata: Asteroidea, representative live images **A** *Ceramaster* gen. inc. (E7048, aboral view) **B** *Ceramaster* gen. inc. (E7048, oral view) **C** *Evoplosoma claguei* (E7081, aboral view) **D** *Evoplosoma claguei* (E7081, oral view) **E** *Nymphaster diomedeeae* (E7235, aboral view) **F** *Nymphaster diomedeeae* (E7235, oral view) **G** *Crossaster borealis* gen. et sp. inc. (E7050, aboral view) **H** *Crossaster borealis* gen. et sp. inc. (E7050, oral view). Scale bars: 1 cm.

Localities. Mound 12 (994–1001 m; this study), Mound 11 (~ 1004–1011 m), Parrita Seep (1097 m; this study).

Distribution. Originally described from 2149 m off Isla del Coco (Ludwig 1905) and recorded from the Gulf of California to Panama and the Galápagos Islands, 702–1618 m (Wehrtmann and Cortés 2009b; Alvarado et al. 2010; Hendrickx et al. 2011; Alvarado and Solís-Marín 2013; Mah 2016).

Echinodermata | Asteroidea | Valvatida | Solasteridae

***Crossaster borealis* Fisher, 1906 gen. et sp. inc.**

Fig. 61G, H

Material examined. AD4917: E7050.

Localities. Mound 12 (1000 m).

Remarks. Possibly *Crossaster borealis*, which was originally described from Alaska and the Bering Sea, 421–1805 m (Fisher 1906) and is known across the northern Pacific from Japan to southern California (Hayashi 1939). If confirmed as *C. borealis*, the CRM specimen would represent a new southern record for the species.

Echinodermata | Asteroidea | Valvatida | Caymanostellidae

***Caymanostella* sp. SIO_BIC_E11440**

Fig. 62A

Material examined. AD4922: MZUCR-ECH2403 (formerly SIO-BIC E11440; [PP627125](#)); AD4972: E7226; AD4976: E7238, E7242.

Localities. Mound 12 (1002 m), Jacó Scar (1845–1887 m).

Remarks. An undescribed species, associated with naturally occurring and experimentally deployed wood, to be described in a separate work (Shen et al. (2024), in press at the time of this work's acceptance).

***Caymanostella* sp. SIO_BIC_E11441**

Fig. 62A

Material examined. AD4503: E4383, E11442; AD4587: E4549, E11221; AD4922: E7075; AD4988: E7289, E7290, E11441 ([PP627110](#)), MZUCR-ECH2402.

Localities. Mound 12 (990–1002 m), Mound 11 (1010 m).

Remarks. An undescribed species, associated with naturally occurring and experimentally deployed wood, to be described in a separate work (Shen et al. (2024), in press at the time of this work's acceptance).

Echinodermata | Ophiuroidea

We list entries following the phylogeny-based higher taxonomy in O'Hara et al. (2018).

Echinodermata | Ophiuroidea | Euryophiurida | Euryalida | Euryalidae

***Asteroschema sublaeve* Lütken & Mortensen, 1899**

Fig. 62B, C

Material examined. AD4916: E7060 (arm fragment only); AD4916: E7061 ([PQ435540](#)); AD4923: E7073 ([PQ435544](#)).

Localities. Parrita Seep (~ 1034–1108 m), Jacó Scar (1604–1710 m).

Distribution. Originally described from several stations in the central eastern Pacific from the vicinity of Islas Marías, Mexico, to Panama, 605–1681 m (Lutken and Mortensen 1899), and also reported from the Gulf of California (Granja-Fernández et al. 2015a).

***Ophiocreas carnosus* Lyman, 1879**

Fig. 62D–F

Material examined. AD4914: E7035; AD4923: E7072 ([PQ435543](#)); S0218: E7318, E7322.

Localities. Parrita Seep (1052 m), Parrita Scar (1431–1589 m), Jacó Scar (1632 m).

Distribution. Originally described from southern Chile (75°S), 320 m (Lyman 1879; Murray and Thomson 1885). Museum records on GBIF include additional occurrences as far north as central Chile (26°S) at depths 175–983 m (GBIF.org 2024).

New records. The CRM specimens represent new northern records and a new maximum depth record for this species (specimen E7035 from 1632 m).

Remarks. Associated with corals: *Callogorgia* (E7072 with MZUCR 3140; E7322 with a similar host, not vouchered), *Swiftia* (E7318 with Co3080), or Isididae (E7035 with MZUCR 3547).

Echinodermata | Ophiuroidea | Euryophiurida | Euryalida |
Gorgonocephalidae

***Gorgonocephalus* cf. *pustulatum* (H.L. Clark, 1916)**

Figs 62G, H, 63A

Material examined. AD4504: E4387 ([PQ435534](#)); AD4922: E7065 ([PQ435542](#)).

Localities. Mound 12 (964 m), Mound 11 (~ 1004–1011 m).

Remarks. Likely an undescribed species, requiring further morphological comparison to the western Pacific *Gorgonocephalus pustulatum*, originally described from the Bass Strait, 183–549 m (Clark 1916b).

***Gorgonocephalus* stet.**

Fig. 63B–D

Material examined. AD4922: E7064 ([PQ435541](#)).

Localities. Mound 12 (965 m).

Echinodermata | Ophiuroidea | Euryophiurida | Ophiurida | Ophiomusina
| Ophiomusaidae

***Ophiomusa lymani* (Wyville Thomson, 1873)**

Fig. 63E–H

Material examined. AD4916: E7045 ([PQ435538](#)); AD4922: E7082 ([PQ435546](#)).

Localities. Mound 12 (~ 966–996 m), Jacó Scar (1674 m).

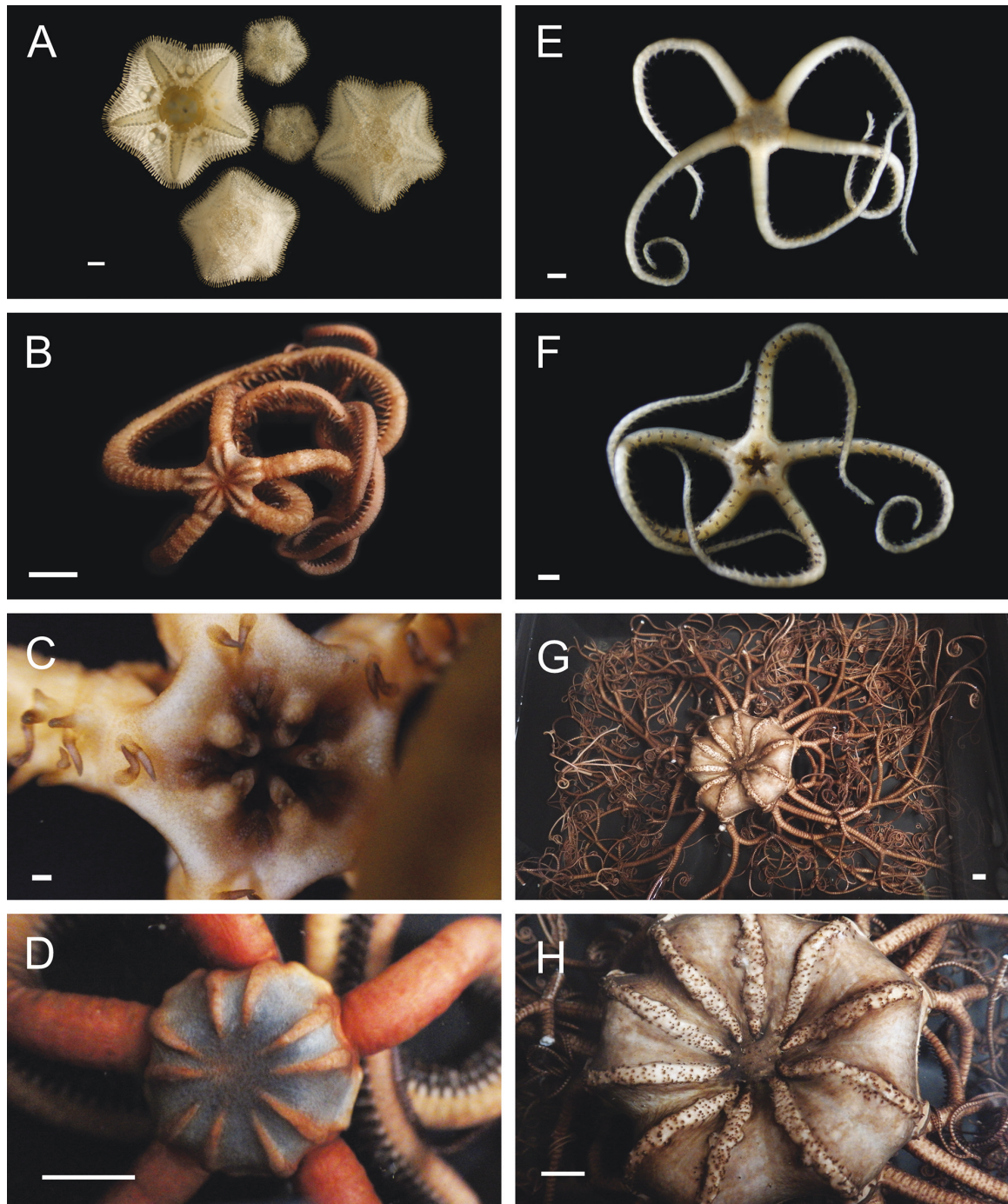


Figure 62. Echinodermata: Asterozoidea and Ophiurozoidea, representative live images **A** *Caymanostella* sp. SIO_BIC_E11440 (E11440) and *Caymanostella* sp. SIO_BIC_E11441 (E7075) as a mixed lot **B** *Asteroschema sublaeve* (E7073, dorsal view) **C** *Asteroschema sublaeve* (E7073, ventral view) **D** *Ophiocreas carnosus* (E7072, dorsal view) **E** *Ophiocreas carnosus* (E7035, dorsal view) **F** *Ophiocreas carnosus* (E7035, ventral view) **G** *Gorgonocephalus* cf. *pustulatum* (E7065, dorsal wide view) **H** *Gorgonocephalus* cf. *pustulatum* (E7065, dorsal view detail). Scale bars: 1 mm (A, C, E, F); 1 cm (B, D, G, H).

Distribution. Originally described from the west coast of Ireland (Thomson 1873) and widely distributed (Baker 2016). *O. lymani* has been reported from the Caribbean and Pacific margins of Central America, including the Galápagos Islands and Isla del Coco, over depths of 51–4700 m (Alvarado et al. 2010; Cortés 2012; Alvarado and Solís-Marín 2013; Granja-Fernández et al. 2023).

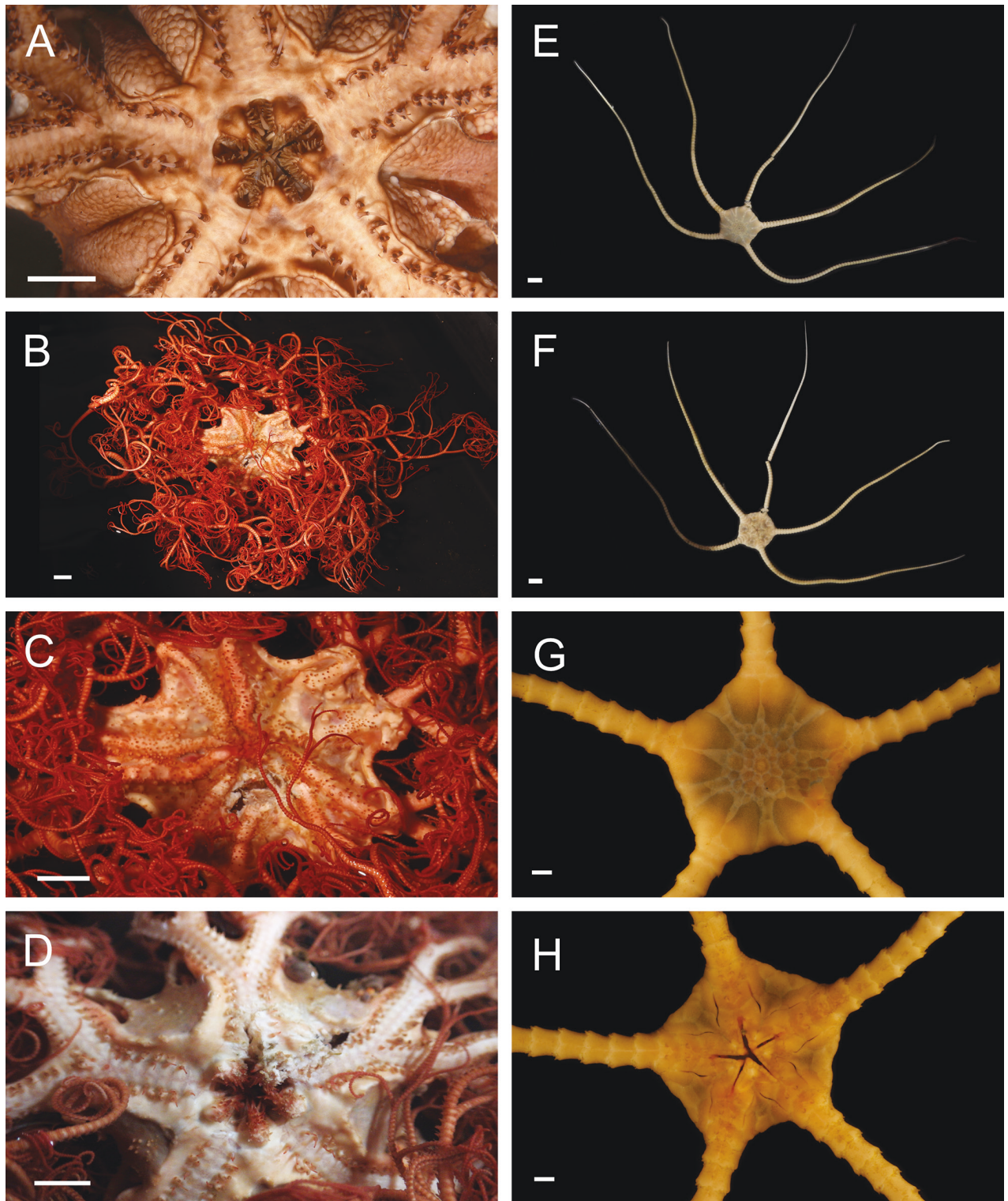


Figure 63. Echinodermata: Ophiuroidea, representative live images **A** *Gorgonocephalus* cf. *pustulatum* (E7065, ventral view detail) **B** *Gorgonocephalus* stet. (E7064, dorsal wide view) **C** *Gorgonocephalus* stet. (E7064, dorsal view detail) **D** *Gorgonocephalus* stet. (E7064, ventral view detail) **E** *Ophiomusa lymani* (E7045, dorsal view) **F** *Ophiomusa lymani* (E7045, ventral view) **G** *Ophiomusa lymani* (E7082, dorsal view detail) **H** *Ophiomusa lymani* (E7082, ventral view detail). Scale bars: 1 cm (A–F); 1 mm (G, H).

Echinodermata | Ophiuroidea | Euryophiurida | Ophiurida | Ophiomusina
| Ophiosphalmidae

***Ophiosphalma glabrum* (Lütken & Mortensen, 1899)**

Fig. 64A, B

Material examined. AD4916: E7046 ([PQ435539](#)).

Localities. Jacó Scar (~ 1854–1887 m).

Distribution. Originally described from several stations in the central eastern Pacific, from the southern Gulf of California to Panama, 1789–4082 m (Lutken and Mortensen 1899), and reported from the Clarion-Clipperton Zone, 4986 m (Christodoulou et al. 2019, 2020). The synonymization of *Ophiomusium multispinum* H.L. Clark, 1911 (described from Washington, 1604 m depth) and *Ophiomusium fimbriatum* Koehler, 1922 (described from the Molucca Sea, Indonesia, 2011 m depth) under *Ophiosphalma glabrum* extends its distribution into the northeastern Pacific and western Pacific, with a total depth range of 878–5203 m (Clark 1913; Baker 2016; Granja-Fernández et al. 2023).

Echinodermata | Ophiuroidea | Euryophiurida | Ophiurida | Ophiurina |
Ophiopyrgidae

***Ophiura flagellata* (Lyman, 1878)**

Fig. 64C, D

Material examined. S0217: E7315.

Localities. The Thumb (1066 m).

Distribution. Originally described from Sagami Bay, Japan, 622 m (Lyman 1878; Murray and Thomson 1885) and widely distributed in the Pacific and Indian Oceans, 96–2330 m (Granja-Fernández et al. 2023). The CRM specimens are morphologically consistent with the synonym *Gymnophiura coerulea* Lütken & Mortensen, 1899, which was originally described from the Mexican Pacific, 1681–1820 m (Lutken and Mortensen 1899), and which likely warrants resurrection (Granja-Fernández et al. 2023).

Remarks. *Ophiura* is polyphyletic with extensive revision required; *O. flagellata* is placed within Ophiopyrgidae (O'Hara et al. 2017).

***Ophiura cf. scutellata* (Lütken & Mortensen, 1899)**

Fig. 64E, F

Material examined. AD4509: E4404; AD4590: E4562 ([PQ450406](#)); AD4912: E7030, E7031, E7032; AD4922: E7066; AD4971: E7230; AD4973: E7231 ([PQ435547](#)); AD4974: E7233; AD4987: E7272.

Localities. Mound 12 (965–1010 m), Jacó Scar (1715–1842 m).

Remarks. Likely an undescribed species, to be further compared to *Ophiura scutellata*, described from Islas Marías, Mexico, 1244 m (Lutken and Mortensen 1899).

***Ophiuroglypha cf. meridionalis* (Lyman, 1879)**

Fig. 64G, H

Material examined. AD4506: E7978; AD4586: E4547 ([PQ435536](#)); AD4587: E4551 ([PQ449362](#)), E4552 ([PQ449363](#)), E7923; AD4907: E7027; AD4977: E7239; AD4985: E7273; S0217: E7321.

Localities. Mound 12 (~ 982–1002 m), Parrita Seep (~ 1030–1033 m), The Thumb (1071 m), Jacó Scar (1783 m).

Remarks. Likely an undescribed species, to be further compared to *Ophiuroglypha meridionalis*, described from the South Atlantic off Río de la Plata, Argentina, 1097 m (Lyman 1879; Murray and Thomson 1885).

***Ophiopyrgidae* sp. SIO_BIC_E7310**

Fig. 65A, B

Material examined. AD4990: E7293; S0214: E7310 ([PQ435550](#)).

Localities. Parrita Seep (1400 m), Jacó Scar (1803 m).

Remarks. An undescribed species, likely also an undescribed genus.

Echinodermata | Ophiuroidea | Ophintegrida | Ophioscolecida |
Ophioscolecidae

***Ophiambix cf. epicopus* Paterson & Baker, 1988**

Fig. 65C, D

Material examined. AD4508: E7979; AD4587: E4550 ([PQ449361](#)); AD4906: E7023; AD4922: E7076; AD4988: E7291.

Localities. Mound 12 (996–1002 m), Mound 11 (1010 m), Parrita Seep (~ 1401–1419 m).

Remarks. Associated with naturally occurring wood falls. Likely an undescribed species, to be further compared to *Ophiambix epicopus*, described from the vicinity of the Kermadec Islands, South Pacific, 530–567 m (Paterson and Baker 1988).

***Ophiophrura liodisca* H.L. Clark, 1911**

Fig. 65E

Material examined. AD4589: E4561; S0219: E7980 ([PQ435551](#)).

Localities. Rio Bongo Scar (606 m), Mound 12 (997 m).

Distribution. Originally described from central Japan, off Omaezaki, 869–924 m (Clark 1911), and recorded from New Zealand, 1421 m (GenBank [KU895351.1](#) (Hugall et al. 2016)). GBIF includes additional occurrences from western Pacific seamounts, 678–1620 m (GBIF.org 2022b).

New records. The CRM specimens represent new records of this species in the eastern Pacific.

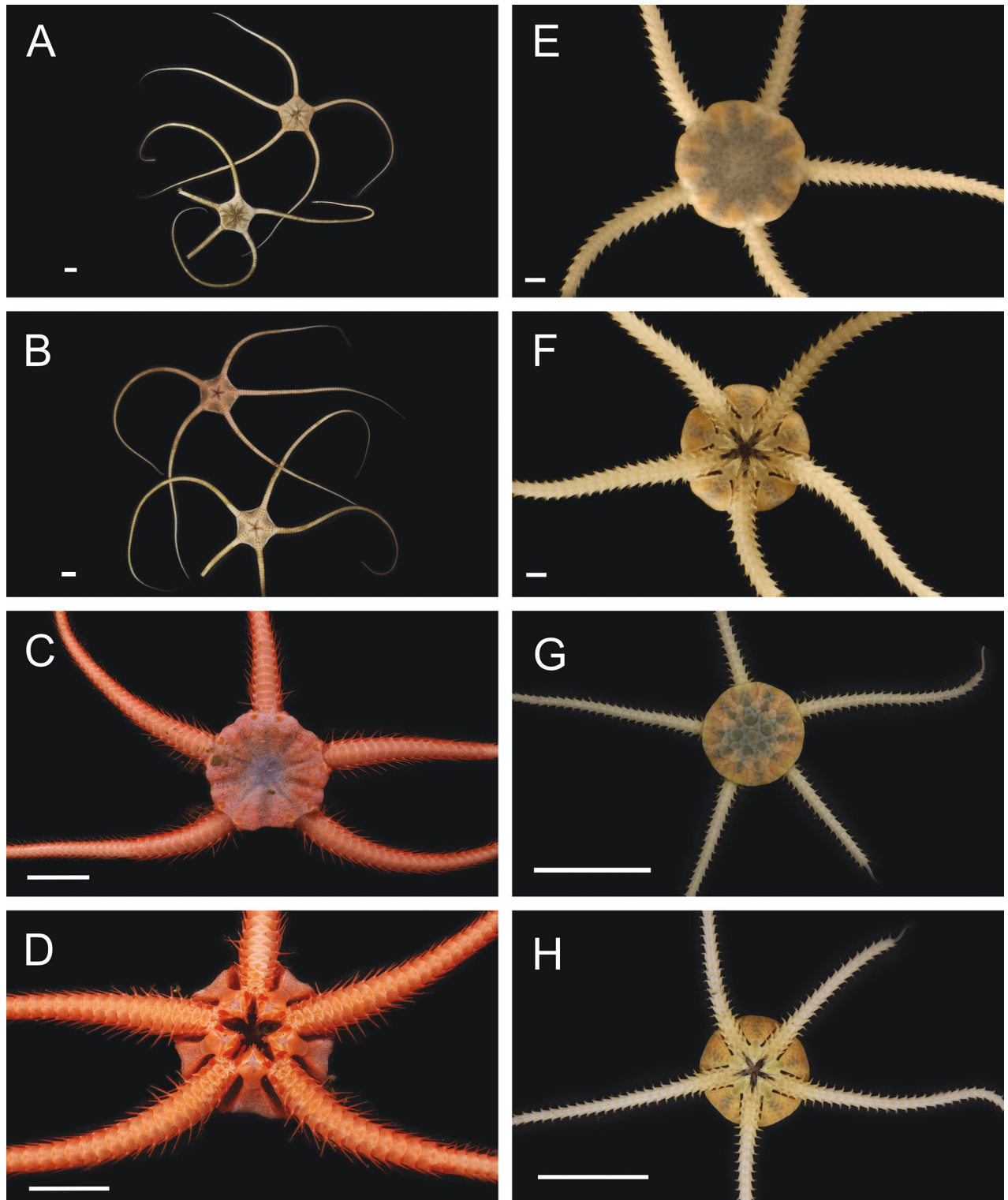


Figure 64. Echinodermata: Ophiuroidea, representative live images **A** *Ophiosphalma glabrum* (E7046, dorsal view) **B** *Ophiosphalma glabrum* (E7046, ventral view) **C** *Ophiura flagellata* (E7315, dorsal view) **D** *Ophiura flagellata* (E7315, ventral view) **E** *Ophiura* cf. *scutellata* (E7231, dorsal view) **F** *Ophiura* cf. *scutellata* (E7231, ventral view) **G** *Ophiuroglypha* cf. *meridionalis* (E7321, dorsal view) **H** *Ophiuroglypha* cf. *meridionalis* (E7321, ventral view). Scale bars: 1 cm (**A–D, G, H**); 1 mm (**E, F**).

Echinodermata | Ophiuroidea | Ophintegrida | Ophiacanthida |
Ophiacanthina | Ophiacanthidae

***Ophiacantha inconspicua* Lütken & Mortensen, 1899**

Fig. 65F

Material examined. AD4913: E7041 ([PQ435552](#)); AD4972: E7245.

Localities. Jacó Scar (~ 1751–1896 m).

Distribution. Originally described and currently known only from several stations off Pacific Panama, 1430–1865 m, and the Galápagos Islands, 1251–1618 m (Lutken and Mortensen 1899).

***Ophiacantha moniliformis* Lütken & Mortensen, 1899**

Fig. 65G, H

Material examined. AD4506: E4390 ([PQ450392](#)); AD4923: E7078; S0219: E7323.

Localities. Rio Bongo Scar (606 m), Parrita Seep (~ 1030–1108 m).

Distribution. Originally described from several stations off Acapulco and Islas Marías, Mexico, 902–1244 m (Lutken and Mortensen 1899) and recorded north to the San Pedro Basin, southern California, 750 m (Hartman 1955; Hartman and Barnard 1958), with an overall reported depth range of 377–1244 m (Granja-Fernández et al. 2023). GBIF includes additional occurrences from Panama and Ecuador, 55–137 m (GBIF.org 2022a).

New records. Pending verification of the GBIF records, our CRM specimens represent new southern records for this species.

***Ophiolebes* sp. SIO_BIC_E4382**

Fig. 66A

Material examined. AD4501: E4382; AD4503: E4385 ([PQ435533](#)).

Localities. Mound 12 (~ 967–995 m).

Remarks. An undescribed species. Associated with the antipatharian coral *Lillipathes ritamariae* (E4382 with Co2293; E4385 with Co2267).

***Ophiolimna* sp. SIO_BIC_E7303**

Fig. 66B, C

Material examined. AD4974: E7234; S0213: E7303 ([PQ435549](#)).

Localities. Jacó Summit (763 m), Mound 12 (992 m).

Remarks. An undescribed species.

Echinodermata | Ophiuroidea | Ophintegrida | Ophiacanthida |
Ophiacanthina | Ophiotomidae

***Ophiomitra partita* Lütken & Mortensen, 1899**

Fig. 66D, E

Material examined. AD4973: E7232 ([PQ435548](#)).

Localities. Jacó Scar (1887 m).

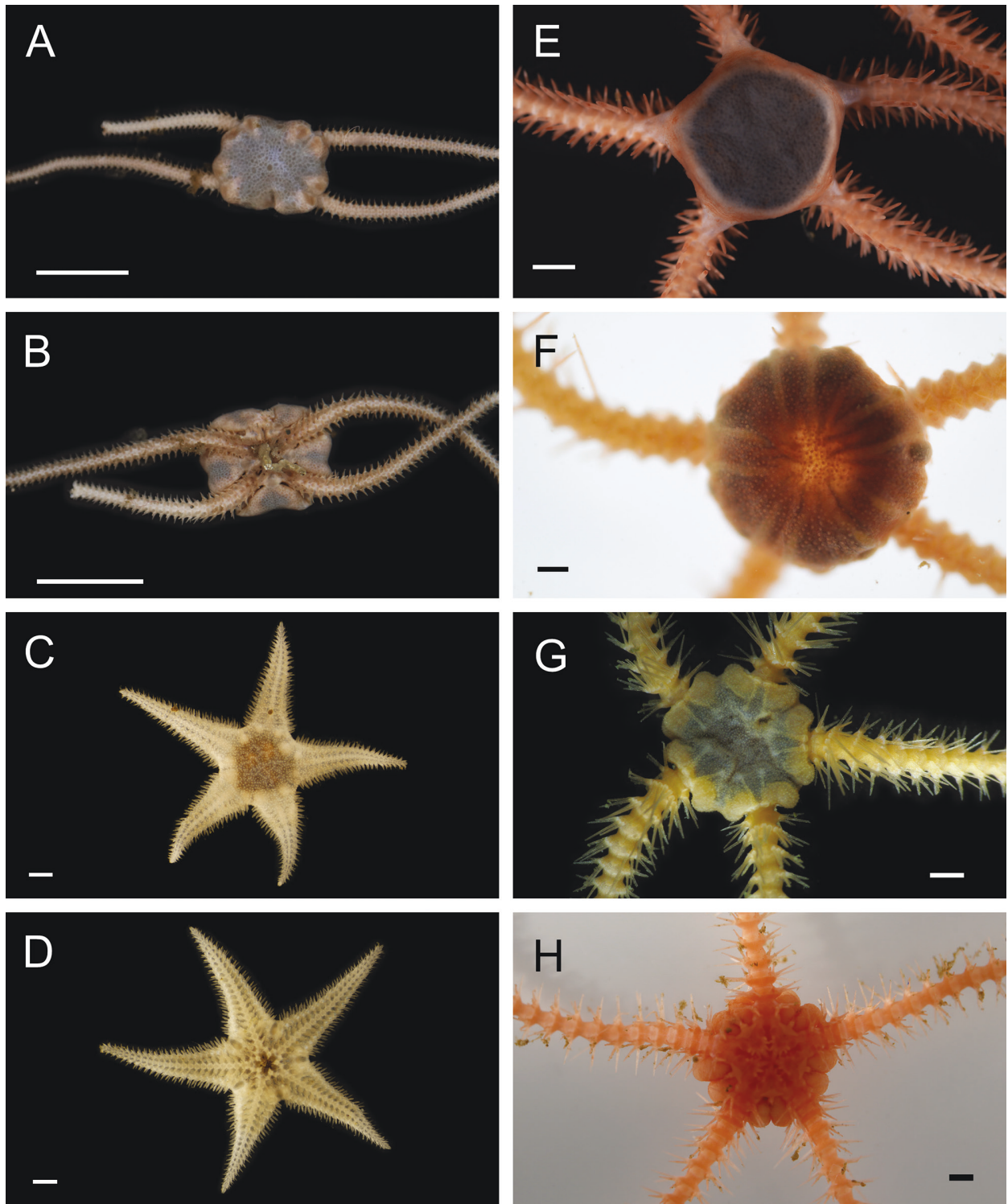


Figure 65. Echinodermata: Ophiuroidea, representative live images **A** Ophiopyrgidae sp. SIO_BIC_E7310 (E7310, dorsal view) **B** Ophiopyrgidae sp. SIO_BIC_E7310 (E7310, ventral view) **C** *Ophiambix* cf. *epicopus* (E7291, dorsal view) **D** *Ophiambix* cf. *epicopus* (E7291, ventral view) **E** *Ophiophrura liodisca* (E4561) **F** *Ophiacantha inconspicua* (E7245) **G** *Ophiacantha moniliformis* (E4390, dorsal view) **H** *Ophiacantha moniliformis* (E7323, ventral view). Scale bars: 1 cm (**A**, **B**); 1 mm (**C**–**H**).

Distribution. Originally described and currently known only from Islas Marías, Mexico, 1236 m (Lutken and Mortensen 1899).

New records. The CRM specimen represents a new southern record and a new maximum depth record for this species.

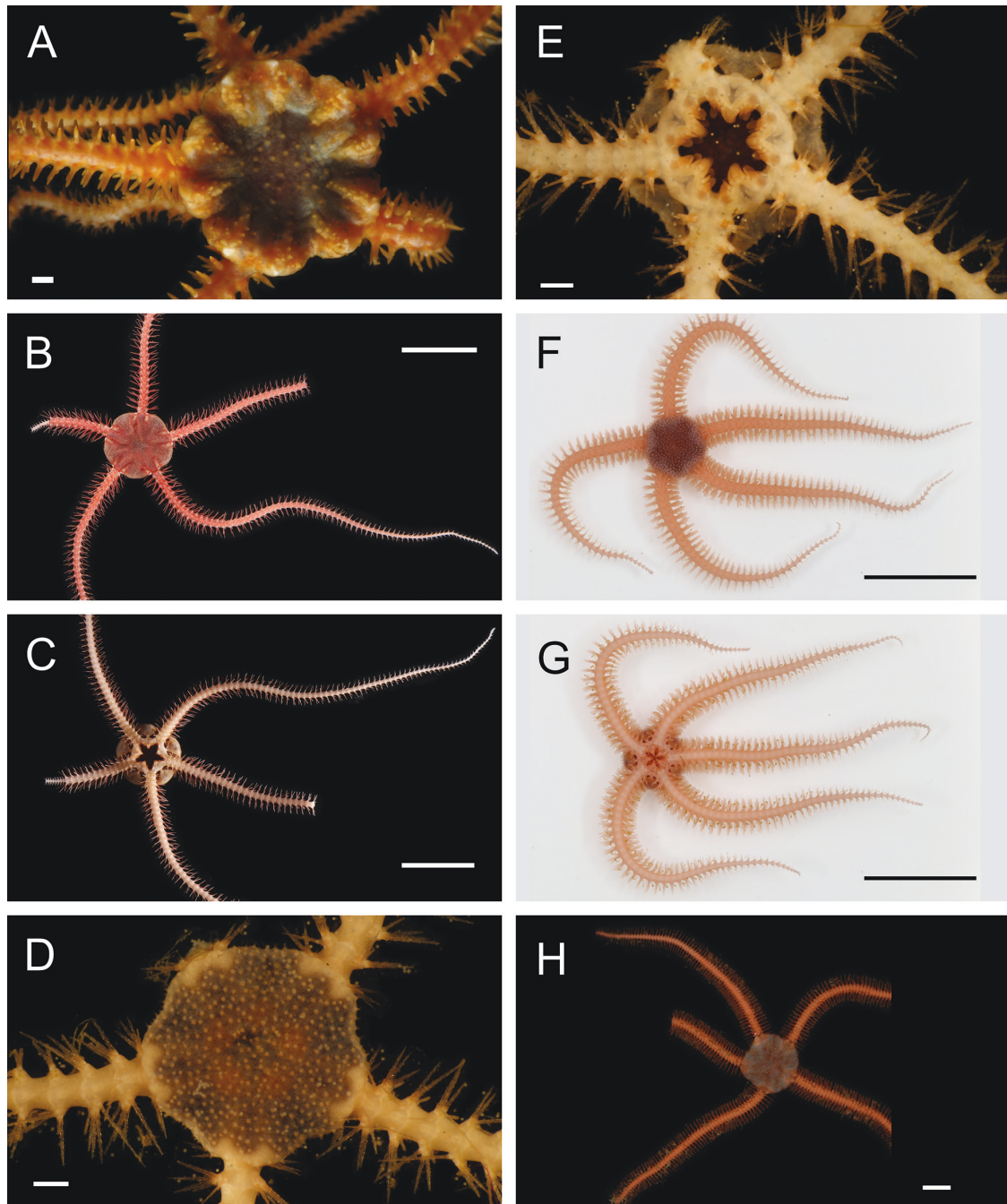


Figure 66. Echinodermata: Ophiuroidea, representative live images **A** *Ophiolebes* sp. SIO_BIC_E4382 (E4382, dorsal view) **B** *Ophiolimna* sp. SIO_BIC_E7303 (E7303, dorsal view) **C** *Ophiolimna* sp. SIO_BIC_E7303 (E7303, ventral view) **D** *Ophiomitra partita* (E7232, dorsal view) **E** *Ophiomitra partita* (E7232, ventral view) **F** *Ophiopristis* sp. SIO_BIC_E7074 (E7074, dorsal view) **G** *Ophiopristis* sp. SIO_BIC_E7074 (E7074, ventral view) **H** *Ophiotreta* sp. SIO_BIC_E4406 (E4406, dorsal view). Scale bars: 1 mm (**A, D, E**); 1 cm (**B, C, F, G**).

***Ophiopristis* sp. SIO_BIC_E7074**

Fig. 66F, G

Material examined. AD4922: E7074 ([PQ435545](#)).

Localities. Mound 12 (~ 967 m).

Remarks. An undescribed species. Associated with the antipatharian coral *Lillipathes ritamariae*.

***Ophiotreta* sp. SIO_BIC_E4406**

Figs 66H, 67A

Material examined. AD4510: E4406 ([PQ435535](#)).

Localities. Jacó Summit (~ 741–744 m).

Remarks. An undescribed species.

Echinodermata | Ophiuroidea | Ophintegrida | Ophioleucida |
Ophioleucidae

***Ophioleuce gracilis* Belyaev & Litvinova, 1976**

Fig. 67B, C

Material examined. AD4913: E7040; AD4916: E7059; AD4971: E7229; AD4972: E7244 (tissue).

Localities. Jacó Scar (1710–1751 m, ~ 1817–1896 m).

Distribution. Known as a bathyal (2000–3000 m) Pacific species (Stöhr et al. 2012) and previously reported from the eastern Clarion-Clipperton Zone, 2882 m (Christodoulou et al. 2019, 2020).

New records. CRM specimen E7059 from 1710 m represents a new minimum depth record for this species.

Echinodermata | Ophiuroidea | Ophintegrida | Amphilepidida |
Gnathophiurina | Amphiuridae

***Amphiura seminuda* Lütken & Mortensen, 1899**

Material examined. AD4587: E7924 ([PQ449044](#); no images available).

Localities. Mound 12 (996 m).

Distribution. Originally described from the southern Gulf of California off Mazatlán, 1558 m (Lutken and Mortensen 1899). Recorded from the San Pedro Basin off southern California, 750 m (Hartman 1955; Hartman and Barnard 1958), to the Gulf of Tehuantepec (Granja-Fernández et al. 2015b), and off the Tuamotu Archipelago, south-central Pacific, 1476 m (Clark 1917).

***Amphiura* sp. SIO_BIC_E4397**

Fig. 67D

Material examined. AD4508: E4397; AD4586: E4563; AD4907: E7033; AD4912: E7029 ([PQ435537](#)).

Localities. Mound 12 (~ 990 m), Jacó Scar (1842 m), Parrita Seep (~ 1400 m).

Remarks. An undescribed species.

Echinodermata | Echinoidea | Echinothurioida | Echinothuriidae

***Araeosoma leptaleum* A. Agassiz & H.L. Clark, 1909**

Fig. 67E–H

Reference. Mongiardino Koch et al. (2018) for phylogenomic analysis.

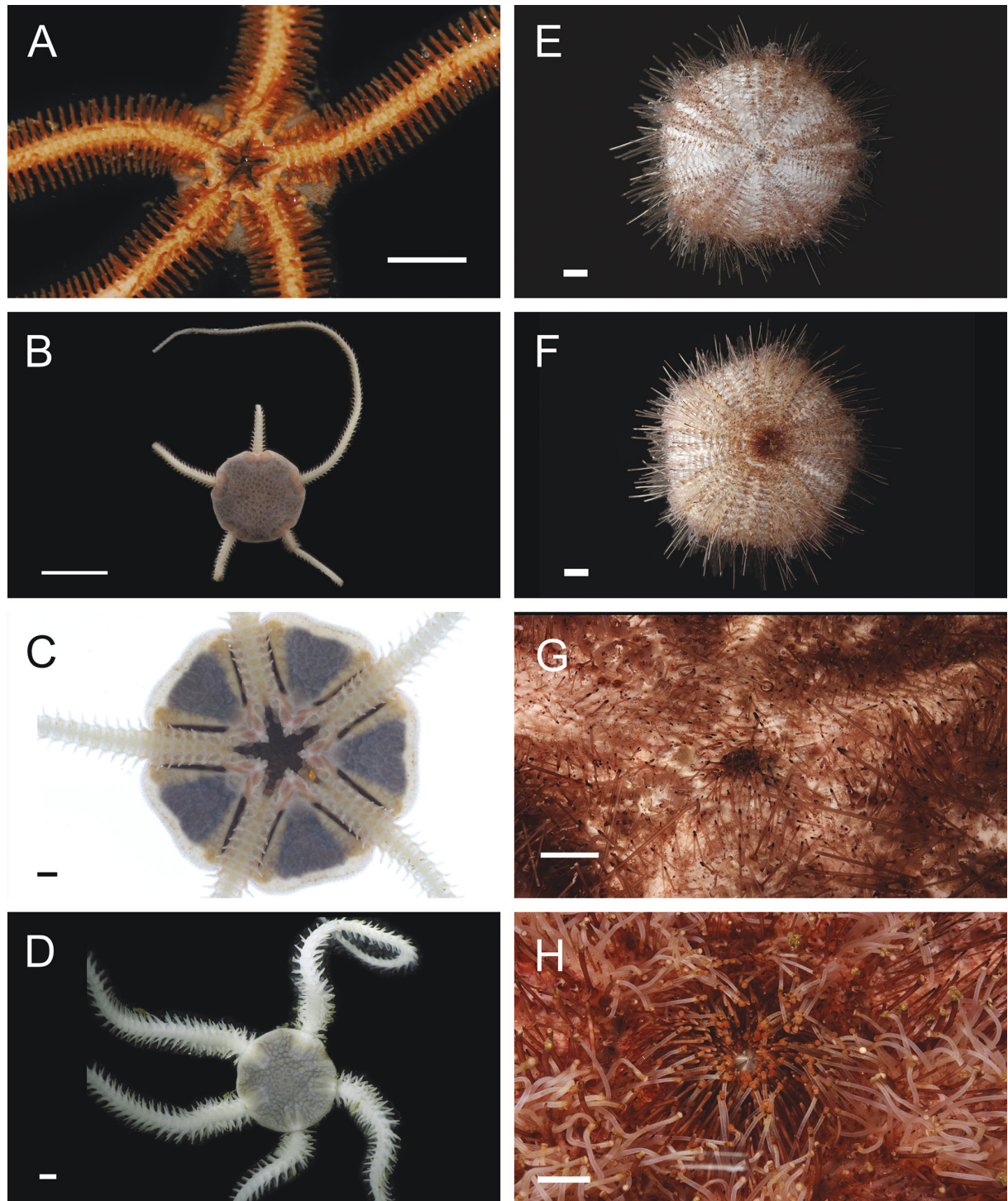


Figure 67. Echinodermata: Ophiuroidea and Echinoidea, representative live images **A** *Ophiotreta* sp. SIO_BIC_E4406 (E4406, ventral view) **B** *Ophioleuce gracilis* (E7040, dorsal view) **C** *Ophioleuce gracilis* (E7244, ventral view) **D** *Amphiura* sp. SIO_BIC_E4397 (E4397, dorsal view) **E** *Araeosoma leptaleum* (E7319, aboral view) **F** *Araeosoma leptaleum* (E7319, oral view) **G** *Araeosoma leptaleum* (E7021, detail of aboral view) **H** *Araeosoma leptaleum* (E7021, detail of oral view). Scale bars: 1 cm (**A**, **B**, **E–H**); 1 mm (**C**, **D**).

Material examined. AD4923: E7021 (transcriptome: SRR7513578); AD4984: E7287; S0218: E7319 ([PQ449387](#)).

Localities. Mound 12 (964 m; this study), Parrita Seep (1091 m), Parrita Scar (1271 m; this study).

Distribution. Known from 740–1063 m along the Pacific margin of Panama (type locality: Mariato Point, 1063 m (Agassiz and Clark 1909)) and the Galápagos seamounts (Mooi et al. 2004; Lessios 2005). GBIF includes occurrences from seamounts off central California, 1093–1106 m (identification confirmed by R. Mooi) (Blum and Fong 2016a, 2016b).

New records. CRM specimen E7319 from 1271 m represents a new maximum depth record for this species.

Echinodermata | Echinoidea | Pedinoidea | Pedinidae

***Caenopedina cf. hawaiiensis* H.L. Clark, 1912**

Fig. 68A, B

Reference. Mongiardino Koch et al. (2018) for phylogenomic analysis.

Material examined. AD4507: E4396 ([PQ450394](#)); AD4913: E7020 (transcriptome: SRR7513589); AD4916: E7047.

Localities. Jacó Scar (1853–1889 m), Parrita Scar (~ 1659–1667 m; this study).

Distribution. *C. hawaiiensis* was originally described from 468–1977 m off the Hawaiian Islands (Clark 1912). GBIF includes occurrences from New Zealand, 392–919 m (Mills and Mackay 2019). Identification of these records and additional records spanning 380–1140 m depth were confirmed by Owen Anderson, National Institute of Water and Atmospheric Research (pers. comm. 22 November 2020). Pending genetic data from the type locality, the specimens from the CRM seeps would represent the first records of this species in the eastern Pacific.

***Caenopedina diomedae* Mortensen, 1939**

Fig. 68C, D

Material examined. AD4984: E7288; S0213: E7304 ([PQ449379](#)).

Localities. Jacó Summit (759 m), Mound 12 (966 m).

Distribution. Originally described from 850 m from the Gulf of Panama (Downey 1968) and reported from 723–933 m in the Gulf of Chiriqui near the Costa Rican border (Lessios 2005).

Echinodermata | Holothuroidea

We list entries following the phylogenetic relationships in Miller et al. (2017) and Mongiardino Koch et al. (2023).

Echinodermata | Holothuroidea | Dendrochirotida | Ypsilothuriidae

***Ypsilothuria bitentaculata* (Ludwig, 1893)**

Fig. 68E

Material examined. AD4509: E4400.

Localities. Jacó Scar (~ 974–1856 m).

Distribution. Originally described from several stations in the eastern Pacific from the Galápagos Islands to central Mexico, 1236–4082 m (Ludwig 1893, 1894), this species is considered circum-Pacific from 135–4000 m, with subspecies

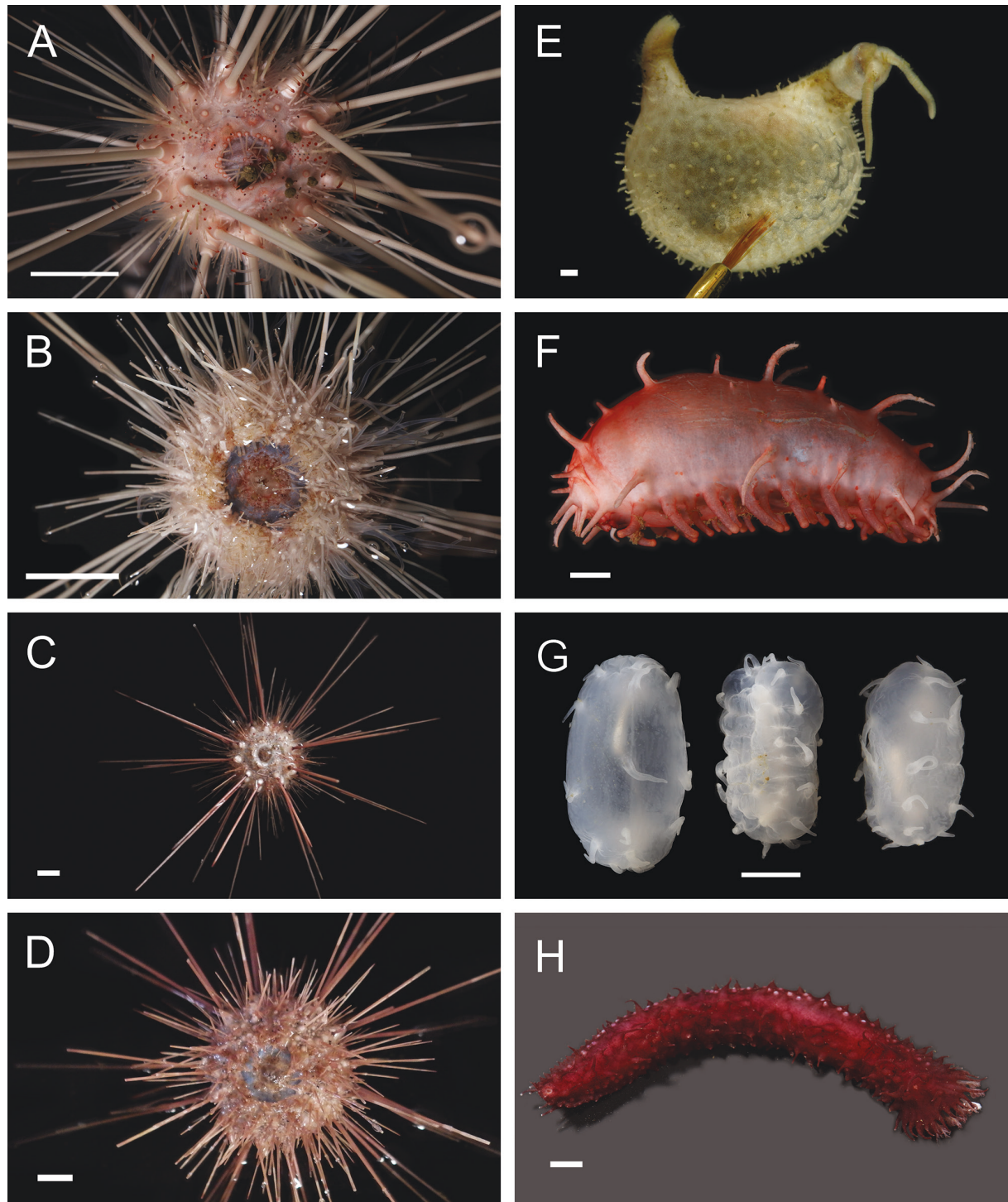


Figure 68. Echinodermata: Echinoidea and Holothuroidea, representative live images **A** *Caenopedina diomedeeae* (E7304, aboral view) **B** *Caenopedina diomedeeae* (E7304, oral view) **C** *Caenopedina* cf. *hawaiiensis* (E7047, aboral view) **D** *Caenopedina* cf. *hawaiiensis* (E7047, oral view) **E** *Ypsilothuria bitentaculata* (E4400) **F** Deimatidae stet. (E7335, adult) **G** Deimatidae stet. (E7336, young) **H** *Bathyplores* cf. *moseleyi* (E7037). Scale bars: 1 cm (**A–D, F–H**); 1 mm (**E**).

or varieties also occurring in the Atlantic and Caribbean (Pawson 1965, 1970; Martins and Tavares 2018). In the eastern Pacific, *Y. bitentaculata* is distributed from central California along the coast of Mexico south to Ecuador (Massin and Hendrickx 2011), including previous reports from Isla del Coco (Wehrtmann and Cortés 2009b) and from 1430 m off northern Panama (Ludwig 1894).

Echinodermata | Holothuroidea | Synallactida | Deimatidae

Deimatidae stet.

Fig. 68F, G

Material examined. S0220: E7335 ([PQ449391](#)), E7336.

Localities. Subduction Plume (3430 m).

Remarks. Female (E7335) with internal brooding of four young (E7336). Likely *Orphnurgus* or *Oneirophanta*.

Echinodermata | Holothuroidea | Synallactida | Synallactidae

***Bathylotes* cf. *moseleyi* (Théel, 1886)**

Figs 68H, 69A

Reference. Mongiardino Koch et al. (2023) for phylogenomic analysis.

Material examined. AD4913: E7037 ([PQ449365](#)), E7038 ([PQ450395](#)); AD4916: E7054 ([PQ449368](#)), E7055 ([PQ449369](#)), E7058 ([PQ449370](#)); S0212: E7302 ([PQ449378](#)); S0214: E7307 ([PQ449380](#)), E7308 ([OR082751](#); transcriptome: SRR24876224), E7309 ([PQ449381](#)).

Localities. Jacó Scar (1847–1875 m).

Remarks. Color variable from deep magenta to brownish to white. Closely related to Antarctic specimens (e.g., SIO-BIC E6356: [KX874357.1](#), COI ~ 96–98% identical) in the *Bathylotes moseleyi* species complex (O’Loughlin et al. 2011). Synallactidae is non-monophyletic and the placement of *Bathylotes* warrants reconsideration (Miller et al. 2017; Mongiardino Koch et al. 2023).

***Bathylotes* sp. SIO_BIC_E7063**

Fig. 69B–D

Material examined. AD4922: E7063 ([PQ449372](#)); S0217: E7314 ([PQ449385](#)).

Localities. Mound 12 (1006 m), The Thumb (1065 m).

Remarks. Color variable, white or pink.

***Synallactes* cf. *chuni* Augustin, 1908**

Fig. 69E–G

Material examined. AD4917: E7049 ([PQ449367](#)); AD4922: E7062 ([PQ449371](#)); S0217: E7313 ([PQ449384](#)).

Localities. Mound 12 (965–966 m), The Thumb (1065 m).

Remarks. E7313 was associated with *Munidopsis aspera* (C13913). These specimens represent the same taxon as voucher specimen E5607, *Synallactes* cf. *chuni*, from the Guaymas Basin, Gulf of California, 1064 m (Miller et al. 2017). *S. chuni* has been reported from the northwestern Pacific, 75–653 m and possibly to 1000 m (Stepanov and Panina 2016).

Echinodermata | Holothuroidea | Elasipodida | Elpidiidae

***Achlyonice* stet.**

Figs 69H, 70A

Material examined. AD4914: E7042, E7043 ([PQ449366](#)); AD4916: E7053, E7056, E7057; S0212: E7297, E7298, E7299, E7300, E7301; S0214: E7305.

Localities. Jacó Scar (~ 1793–1869 m).

Remarks. Three specimens (E7299, E7305, E7306) were collected swimming 1–9 m above the seafloor; the other specimens were benthic. The closest COI BLASTN result on GenBank was an unidentified elpidiid from the Indian Ocean ([PP778426.1](#), voucher NMV F296840, 93.53% identity).

Echinodermata | Holothuroidea | Elasipodida | Laetmogonidae

We thank Akito Ogawa (Japan Agency for Marine-Earth Science and Technology) for input on these identifications.

Laetmogonidae fam. inc.

Fig. 70B

Material examined. AD4923: E7068 ([PQ449373](#)), E7071 ([PQ449375](#)).

Localities. Parrita Seep (1108 m).

Distribution. These specimens morphologically resemble *Laetmogone scoeoides* (H.L. Clark, 1913), which was described from Ballenas Bay, Baja California, 1180 m (Clark 1913), and reported to 1420 m off Guerrero, Mexico (Granja-Fernández et al. 2015b), but further comparison to the type material is required (Akito Ogawa, pers. comm. 10 August 2022). The COI sequences did not closely match any available GenBank reference sequences (<84% identity), and phylogenetic interpretation is challenging, as Laetmogonidae is paraphyletic (Miller et al. 2017; Mongiardino Koch et al. 2023).

***Pannychia* sp. SIO_BIC_E7080**

Fig. 70C, D

Material examined. AD4923: E7080 ([PQ450401](#)); S0218: E7317 ([PQ449386](#)).

Localities. Parrita Seep (~ 1037–1108 m), Parrita Scar (1364 m).

Remarks. A potentially undescribed species, within the *Pannychia moseleyi* Théel, 1882 complex, which requires extensive revision (Ogawa et al. 2022, 2023).

***Pannychia* sp. SIO_BIC_E7286**

Fig. 70E

Material examined. AD4990: E7286 ([PQ449377](#)).

Localities. Parrita Seep (1408 m).

Remarks. A potentially undescribed species, within the *Pannychia moseleyi* complex, which requires extensive revision (Ogawa et al. 2022, 2023).

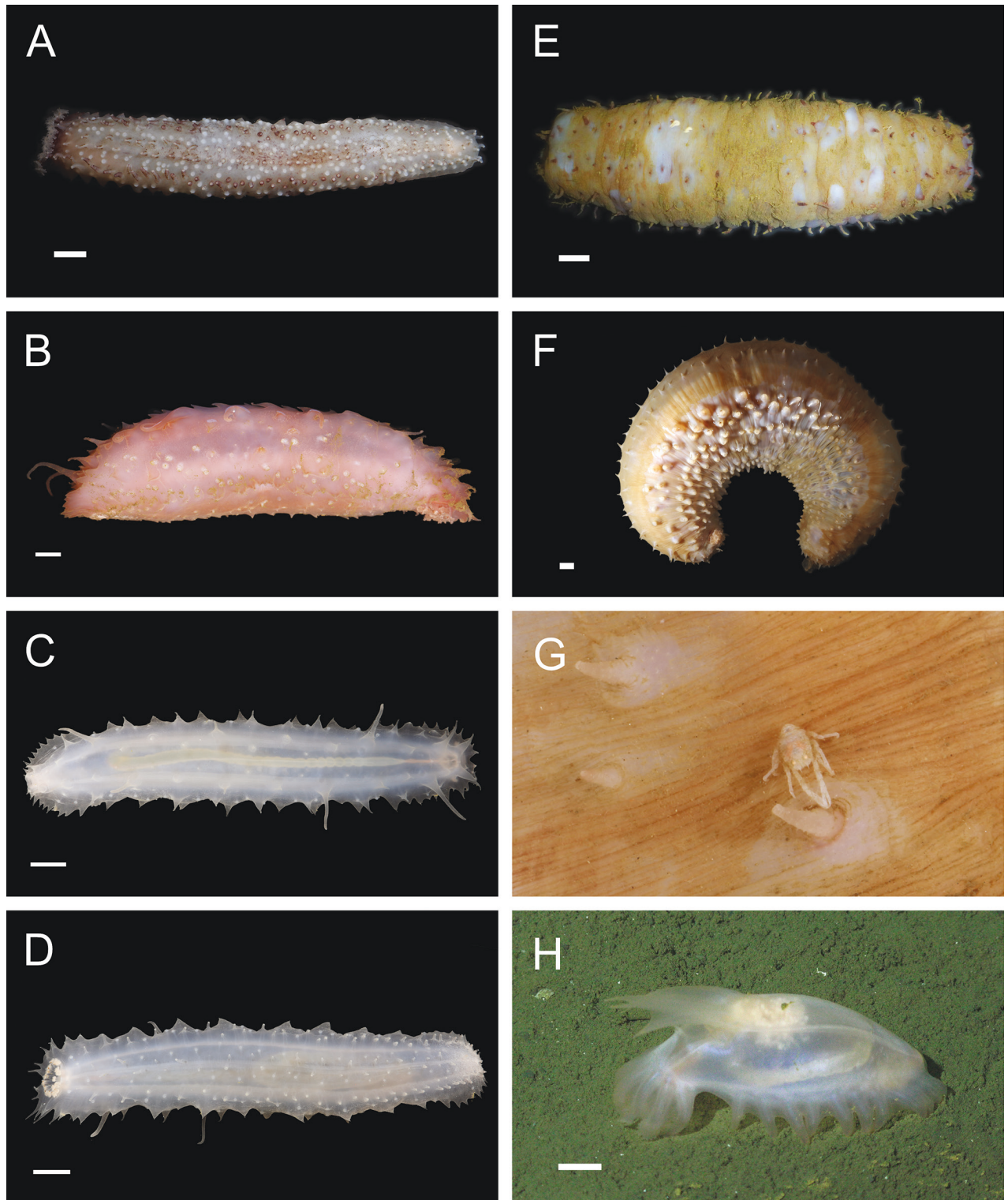


Figure 69. Echinodermata: Holothuroidea, representative live images **A** *Bathyplores* cf. *moseleyi* (E7038) **B** *Bathyplores* sp. SIO_BIC_E7063 (E7063) **C** *Bathyplores* sp. SIO_BIC_E7063 (E7314, dorsal view) **D** *Bathyplores* sp. SIO_BIC_E7063 (E7314, ventral view) **E** *Synallactes* cf. *chuni* (E7049, dorsal view) **F** *Synallactes* cf. *chuni* (E7313, lateral view) **G** *Synallactes* cf. *chuni* (E7313, detail with *Munidopsis aspera*) **H** *Achlyonice* stet. (E7297, *in situ*). Credit: ROV SuBastian/Schmidt Ocean Institute. Scale bars: 1 cm.

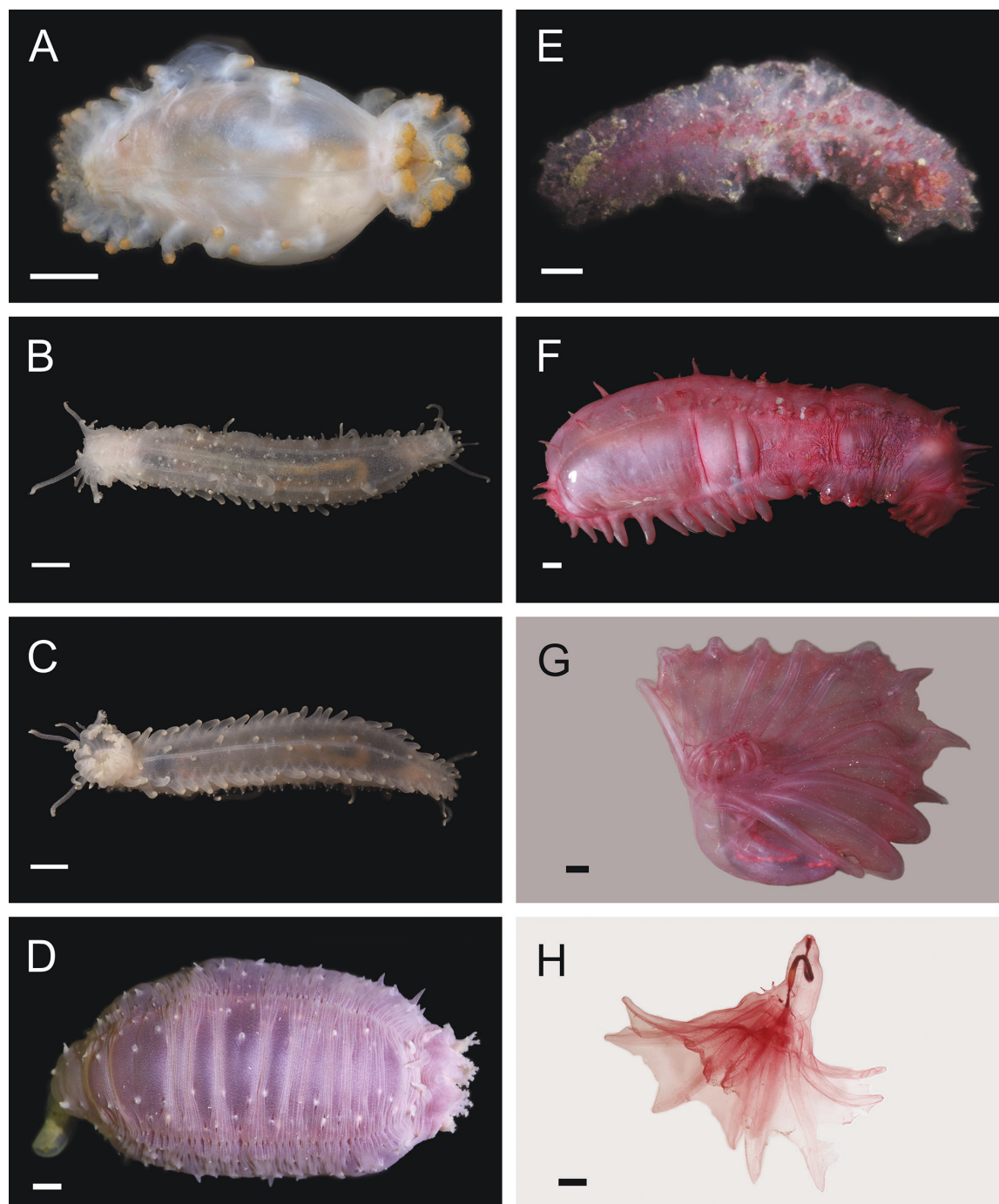


Figure 70. Echinodermata: Holothuroidea, representative live images **A** *Achlyonice* stet. (E7042, ventral view) **B** *Pannychia* sp. SIO_BIC_E7080 (E7317, dorsal view) **C** *Pannychia* sp. SIO_BIC_E7080 (E7317, ventral view) **D** *Pannychia* sp. SIO_BIC_E7286 (E7286) **E** *Pannychia* stet. (E7070) **F** Laetmogonidae fam. inc. (E7068) **G** *Pelagothuria natatrix* (E7311, frontal view) **H** *Pelagothuria natatrix* (E7312, lateral view). Scale bars: 1 cm.

***Pannychia* stet.**

Fig. 70F

Material examined. AD4923: E7070 ([PQ449374](#)), E7079 ([PQ449376](#)).

Localities. Parrita Seep (1091–1098 m).

Remarks. A potentially undescribed species, near the *Pannychia moseleyi* complex.

Echinodermata | Holothuroidea | Elasipodida | Pelagothuriidae

***Pelagothuria natatrix* Ludwig, 1893**

Fig. 70G, H

Material examined. S0215: E7311 ([PQ449382](#)); S0217: E7312 ([PQ449383](#)).

Localities. Mound 12 (982 m depth, 1 m above the seafloor), The Thumb (1056 m depth, 11 m above the seafloor).

Distribution. Originally described from several stations in the eastern Pacific from the Gulf of Panama to the Galápagos Islands, 605–3350 m (Ludwig 1893, 1894) and previously reported from Isla del Coco (Cortés 2012). Globally, *Pelagothuria* has been reported from the tropical regions of all ocean basins, at depths of 197–4441 m, but collection of specimens is important to ascertain whether these records represent a single species (Selig et al. 2019).

Echinodermata | Holothuroidea | Elasipodida | Psychropotidae

***Benthodytes* cf. *sanguinolenta* Théel, 1882**

Fig. 71A

Material examined. S0220: E7334 ([PQ449390](#)).

Localities. Subduction Plume (3453 m depth, 2 m above the seafloor).

Remarks. Collected while swimming. The closest COI BLASTN results on GenBank were *Benthodytes* cf. *sanguinolenta* from the Indian Ocean ([PP778424.1](#), voucher NMV F308225, 98.91% identity) and an Antarctic specimen of *B. sanguinolenta* ([HM196505.1](#), 97.85% identity), which is a species complex (O’Loughlin et al. 2011).

Echinodermata | Crinoidea

We thank and commemorate Charles Messing (Nova Southeastern University) for assistance with these identifications.

Echinodermata | Crinoidea | Comatulida | Antedonoidea | Antedonidae

***Fariometra* sp. SIO_BIC_E4389**

Fig. 71B

Material examined. AD4506: E4389 ([PQ449360](#)).

Localities. Parrita Seep (~ 1030–1179 m).

Remarks. Antedonidae is not monophyletic and requires major revision (Hemery et al. 2013). This specimen warrants comparison to *Fariometra parvula* (Hartlaub, 1895) which has been recorded from the tropical eastern Pacific including Costa Rica (Alvarado et al. 2022).

***Fariometra* sp. SIO_BIC_E4399**

Fig. 71C, D

Reference. Summers et al. (2014a) for DNA sequences (KM014345, reported as *Antedon* sp.) and phylogenetic analysis of specimen E4399.

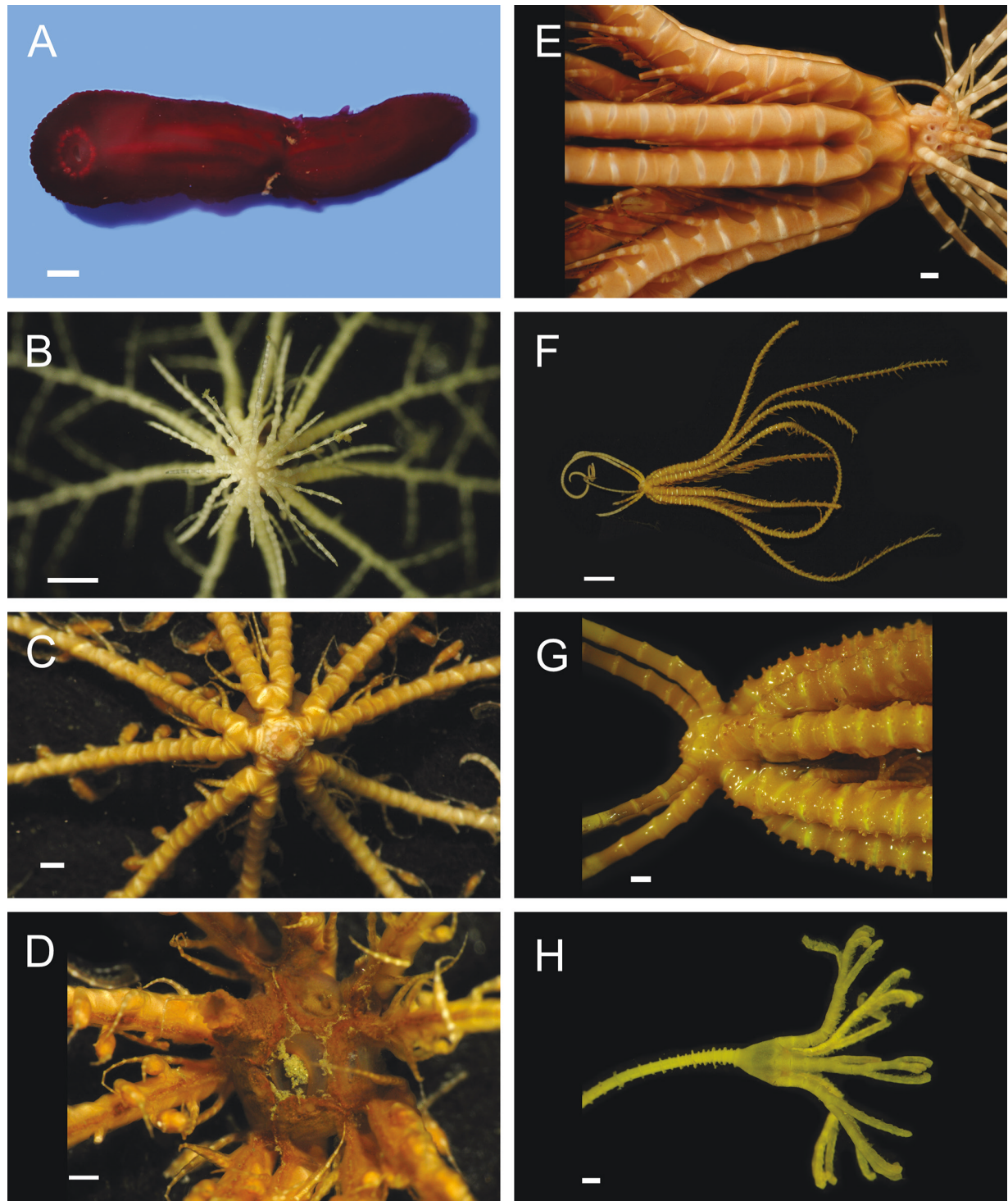


Figure 71. Echinodermata: Holothuroidea and Crinoidea, representative live images **A** *Benthodytes* cf. *sanguinolenta* (E7334) **B** *Fariometra* sp. SIO_BIC_E4389 (E4389) **C** *Fariometra* sp. SIO_BIC_E4399 (E4399, aboral view) **D** *Fariometra* sp. SIO_BIC_E4399 (E4399, oral view) **E** *Psathyrometra* cf. *fragilis* (E7034) **F** *Thalassometra agassizii* (E4391, lateral view) **G** *Thalassometra agassizii* (E4391, detail) **H** *Calamocrinus diomedae* (E4394). Scale bars: 1 cm (**A**, **F**); 1 mm (**B–E**, **G**, **H**).

Material examined. AD4509: E4399.

Localities. Jacó Scar (1789 m).

Remarks. Host of *Pulvinomyzostomum inaki* (A1408 and A1579) (Summers et al. 2014b). This specimen represents a different morphospecies than E4389, with COI 96.15% identical, also warranting comparison to *Fariometra parvula*.

Echinodermata | Crinoidea | Comatulida | Antedonoidea | Zenometridae

***Psathyrometra cf. fragilis* (AH Clark, 1907)**

Fig. 71E

Material examined. AD4913: E7034 ([PQ449364](#)).

Localities. Jacó Scar (1878 m).

Distribution. *Psathyrometra fragilis* has been reported along the Pacific Rim from Japan (type locality, 914 m) (Clark 1907) through the Aleutian Islands and California to the Gulf of Panama, 439–2903 m (Messing and White 2001).

Remarks. Host of *Myzostoma josefinae* (A8362) and *Pulvinomyzostomum* sp. SIO_BIC_A8361.

Echinodermata | Crinoidea | Comatulida | Tropiometroidea |
Thalassometridae

***Thalassometra agassizii* (Hartlaub, 1895)**

Fig. 71F, G

Material examined. AD4506: E4391 ([PQ450400](#)).

Localities. Parrita Seep (~ 1030–1179 m).

Distribution. Originally described from the Galápagos Islands, 598–1250 m, and Panama, 1429 m (Clark 1915) and previously reported from Isla del Coco (Wehrtmann and Cortés 2009b).

Echinodermata | Crinoidea | Hyocrinida | Hyocrinidae

***Calamocrinus diomedae* Agassiz, 1890**

Fig. 71H

Material examined. AD4507: E4394 ([PQ450393](#)).

Localities. Parrita Scar (~ 1659–1667 m).

Distribution. Originally described from 717 m off the Galápagos Islands (Agassiz 1890) and known from Pacific Panama and Isla del Coco over a total depth range of 525–1430 m (Roux 2004).

New records. The CRM specimen represents a new depth record for this species (1659 m as the most conservative value).

Hemichordata

***Saccoglossus* sp. SIO_BIC_H35**

Fig. 72A

Material examined. AD4906: H35 (16S: [PQ304659](#)).

Localities. Mound 12 (1000 m).

Remarks. The closest 16S BLASTN result on GenBank was a specimen of *Saccoglossus* sp. from Friday Harbor, Washington, 5–10 m ([KF683544.1](#), 96.40% identity). Based on the reported 16S interspecific distances within the monophyletic genus *Saccoglossus* (0.1–17.9%, corrected) and distances

between *Saccoglossus* and other harrimaniid genera (>18.2%, corrected) (Cannon et al. 2013), we identify the CRM specimen as within *Saccoglossus* and likely an undescribed species.

Xenacoelomorpha

Xenoturbella sp. SIO_BIC_BI1373

Fig. 72B–D

Material examined. S0215: BI1364; S0230: BI1372, BI1373, BI1374.

Localities. Mound 12 (999 m), Mound Jaguar (1895 m).

Remarks. An undescribed species.

Cnidaria

Cnidaria | Anthozoa

We list entries following the phylogenetic relationships in McFadden et al. (2021).

Cnidaria | Anthozoa | Hexacorallia | Actiniaria | Actinioidea |
Liponematidae

Liponema stet.

Fig. 72E

Material examined. AD4913: Co2922.

Localities. Jacó Scar (1871 m).

Cnidaria | Anthozoa | Hexacorallia | Actiniaria | Metridioidea |
Hormathiidae

Hormathiidae stet.

Fig. 72F, G

Material examined. AD4504: Co2268; AD4513: Co2280 (16S: [PQ304656](#); COIII: [PQ435166](#)); AD4907: Co2918; AD4922: Co2875, Co2933; AD4923: Co2877.

Localities. Mound 11 (~ 1004–1011 m), Mound 12 (994–1002 m), Parrita Seep (1101 m), Jacó Scar (~ 1799 m).

Remarks. Multiple species may be represented. Co2268 was attached to the tube of *Lamellibrachia donwalshi*. Co2875 and Co2933 were associated with squat lobsters, *Munidopsis hendersoniana* (C12803 and C12799, respectively). For Co2280 the closest BLASTN results on GenBank were: for 16S, 99.78% identity to *Paraphelliactis xishaensis* Feng, Liu, Xu, Zhou, Zhu, Liu, Wu, Li, Qiu, He, Wang, Zhang & Wang, 2021 ([MT997141.1](#)), *Paracalliactis* sp. ([FJ489429.1](#)), and *Paraphelliactis* sp. ([FJ489431.1](#)); for COIII, 100.00% identity to unidentified Hormathiidae from central California at 1287 m ([MN954949.1](#)) and *Chondrophellia* sp. ([FJ489489.1](#)).

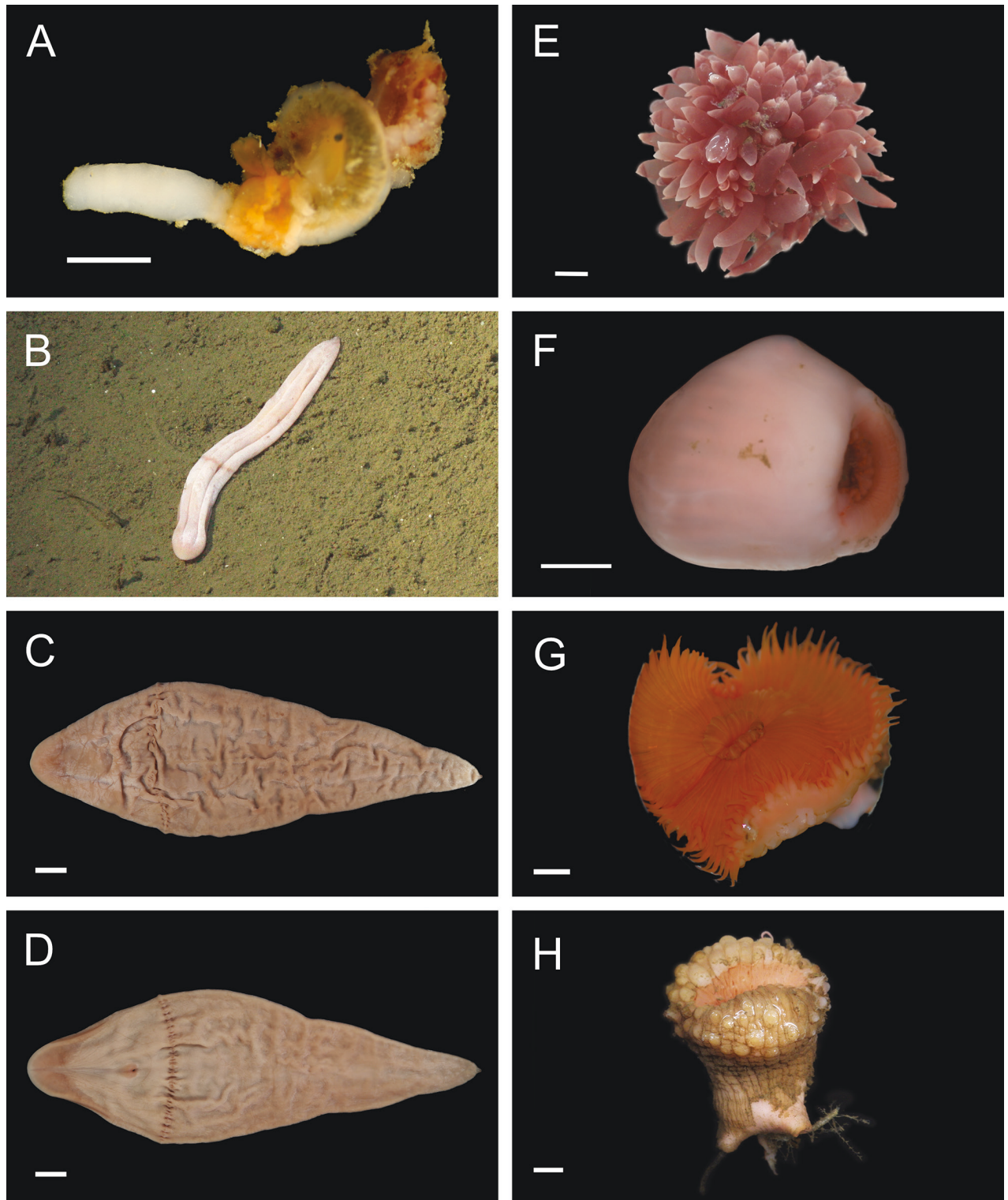


Figure 72. Hemichordata, Xenacoelomorpha, and Cnidaria: Actiniaria, representative live images **A** *Saccoglossus* sp. SIO_BIC_H35 (H35) **B** *Xenoturbella* sp. SIO_BIC_BI1373 (BI1364, *in situ*). Credit: ROV SuBastian/Schmidt Ocean Institute **C** *Xenoturbella* sp. SIO_BIC_BI1373 (BI1373, dorsal view) **D** *Xenoturbella* sp. SIO_BIC_BI1373 (BI1373, ventral view) **E** *Liponema* stet. (Co2922) **F** *Paracalliactis* stet. (Co2300) **G** Hormathiidae stet. (Co2268) **H** Hormathiidae stet. (Co2875). Scale bars: 1 mm (**A**); 1 cm (**C–H**).

***Paracalliactis* stet.**

Fig. 72H

Material examined. AD4590: Co2300; AD4591: Co2320.

Localities. Jacó Scar (~ 1800 m).

Remarks. Associated with *Parapagurus foraminosus* hermit crabs, as is characteristic of *Paracalliactis* (Gusmão and Daly 2010; Gusmão et al. 2020) (Co2300 with crab C11210, Co2320 with crab C13308).

Cnidaria | Anthozoa | Hexacorallia | Actiniaria | Metridioidea |
Kadosactinidae

Kadosactinidae cf. sp. B sec. Goffredi et al. 2021

Fig. 73A, B

Material examined. AD4501: Co2265 (16S: [PQ304653](#); COIII: [PQ299038](#)); AD4513: Co2279 (16S: [PQ304655](#)); AD4590: Co2309 ([PQ449359](#); 16S: [PQ304657](#)).

Localities. Mound 12 (~ 984–997 m), Jacó Scar (~ 1744–1818 m).

Remarks. Co2309 was associated with a vestimentiferan tubeworm (*Escarpia spicata* or *Lamellibrachia barhami*). The COIII sequence of Co2265 was nearly identical to those of an undescribed vent anemone, Kadosactinidae sp. B, from the Pescadero Basin, Gulf of California, 3692 m ([MW148236.1](#), 99.46% identity) (Goffredi et al. 2021), and to *Alvinactis chessi* Zelnio, Rodríguez & Daly, 2009 (Kadosactinidae) ([GU473352.1](#), 99.17% identity). Due to the relatively short sequence lengths and close relationships among taxa, the CRM 16S sequences were 100.0% identical to those of Kadosactinidae sp. B ([MW172213.1](#)), *A. chessi* ([GU473296.1](#)), and *Cyananthea hourdezi* Zelnio, Rodríguez & Daly, 2009 (Kadosactinidae) ([GU473293.1](#)). The COI sequence of Co2309 was 99.52–99.68% identical to those of Actinostolidae spp. from vents in the Indian Ocean ([MH202753.1](#), [OK267405.1](#), [OK267413.1](#)) and *Maractis* sp. from vents at the Mid-Cayman Spreading Center ([KJ566948.1](#)), although no COI sequences from any Kadosactinidae were available for comparison.

Cnidaria | Anthozoa | Hexacorallia | Actiniaria | Metridioidea

Amphianthidae fam. inc.

Fig. 73C, D

Material examined. AD4506: Co2273 (16S: [PQ304654](#); COIII: [PQ299039](#)); AD4923: Co2874.

Localities. Parrita Seep (~ 1030–1094 m).

Remarks. Associated with coralliids (Co2273 with coralliid Co2271; Co2874 with coralliid Co2947). For Co2273 the closest BLASTN results on GenBank were: for 16S, *Peronanthus* sp. ([KJ482956.1](#), 99.56% identity), *Galatheanthemum* sp. from the Mariana Trench ([OL912950.1](#), 99.33% identity), and an unidentified sea anemone (U40291.1, 99.33% identity); for COIII, 95.94% identity to *Peronanthus* sp. ([KJ482976.1](#)) and *Galatheanthemum* spp. ([KJ482977.1](#), [OL912950.1](#), [OQ697433.1](#)).

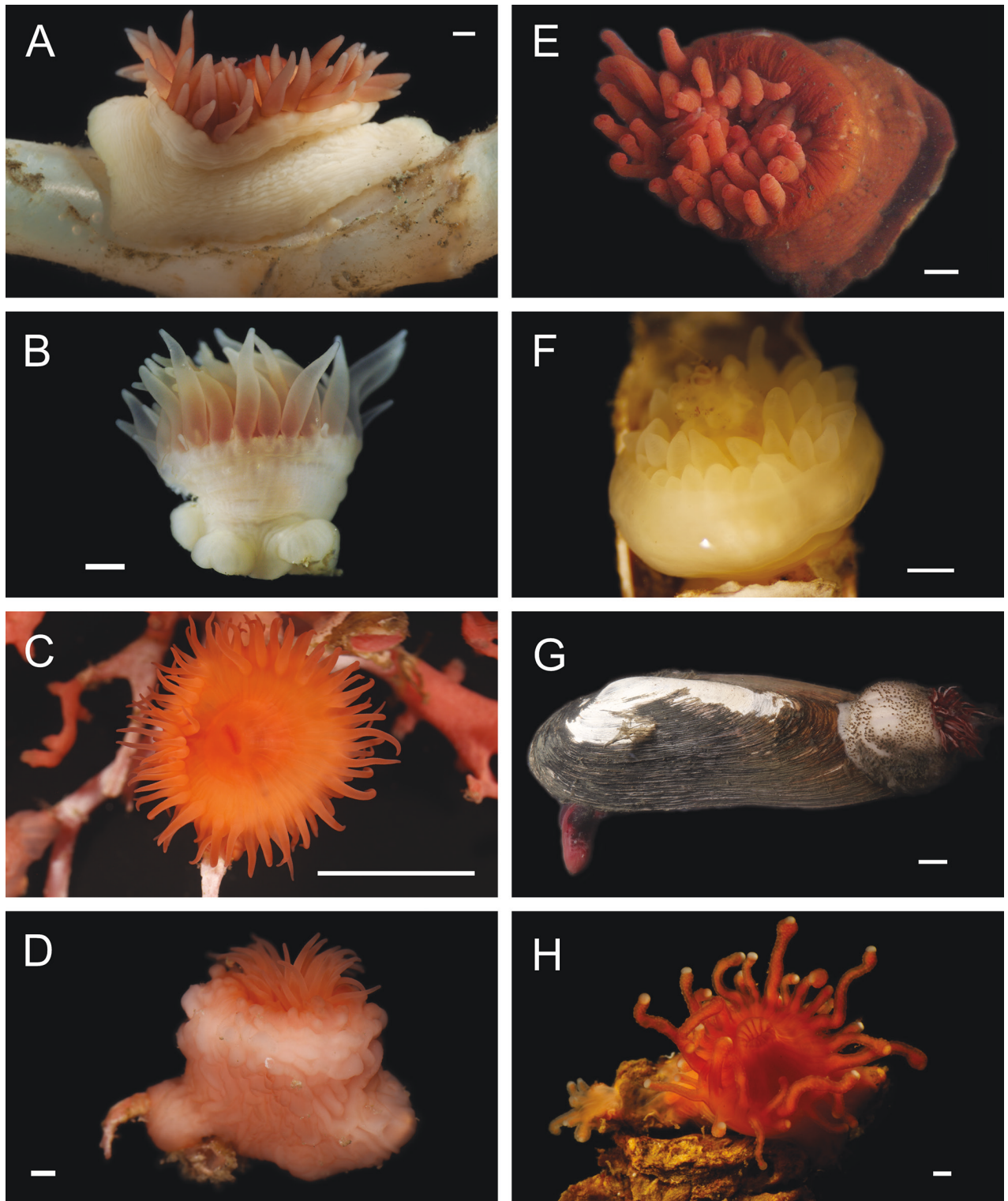


Figure 73. Cnidaria: Actiniaria, representative live images **A** Kadosactinidae cf. sp. B sec. Goffredi et al. 2021 (Co2309, oral view) **B** Kadosactinidae cf. sp. B sec. Goffredi et al. 2021 (Co2265, lateral view) **C** Amphianthidae fam. inc. (Co2874, oral view) **D** Amphianthidae fam. inc. (Co2874, lateral view) **E** Actiniaria sp. SIO_BIC_Co2876 (Co2876) **F** Actiniaria sp. SIO_BIC_Co3056 (Co3056) **G** Actiniaria sp. SIO_BIC_Co3084 (Co3084) **H** Actiniaria sp. SIO_BIC_Co3086 (Co3086). Scale bars: 1 mm (**A, B, D, F, H**); 1 cm (**C, E, G**).

Cnidaria | Anthozoa | Hexacorallia | Actiniaria

Actiniaria sp. SIO_BIC_Co2876

Fig. 73E

Material examined. AD4923: Co2876.

Localities. Parrita Seep (1041 m).

Actiniaria sp. SIO_BIC_Co3056

Fig. 73F

Material examined. AD4988: Co3056.

Localities. Mound 11 (1010 m).

Remarks. Associated with a naturally occurring wood fall.

Actiniaria sp. SIO_BIC_Co3084

Fig. 73G

Material examined. S0220: Co3084.

Localities. Subduction Plume (3408 m).

Remarks. Associated with a clam, *Calyptogena diagonalis* (M17065).

Actiniaria sp. SIO_BIC_Co3086

Fig. 73H

Material examined. S0219: Co3086.

Localities. Rio Bongo Scar (661 m).

Remarks. Associated with a naturally occurring wood fall.

Cnidaria | Anthozoa | Hexacorallia | Antipatharia | Schizopathidae

***Bathypathes* stet.**

Fig. 74A, B

Material examined. AD4914: MZUCR 3537; AD4922: MZUCR 3541; S0218: MZUCR 3545.

Localities. Mound 12 (966 m), Parrita Scar (1154 m), Jacó Scar (1632 m).

***Lillipathes ritamariae* Opresko & Breedy, 2010**

Fig. 74C

Reference. Opresko and Breedy, 2010** (no DNA sequences available).

Localities. Mound 12 (~ 967–995 m; type locality).

Distribution. Recorded in the eastern Pacific from the CRM and northern Chile, and in the southwestern Pacific off New Zealand and Antarctica, 1005–1746 m (Araya et al. 2018).

Remarks. Host of the myzostome *Eenymeenymyzostoma* sp. SIO_BIC_A8428.

Cnidaria | Anthozoa | Hexacorallia | Scleractinia | Flabellidae

***Javania cailleti* (Duchassaing & Michelotti, 1864)**

Fig. 74D

Material examined. AD4510: Co2276.

Localities. Jacó Summit (~ 741–744 m).

Distribution. Originally described from the Caribbean (Duchassaing and Michelotti 1864) and considered cosmopolitan (Cairns 1991). Previously reported from Isla del Coco and the Galápagos Islands, 245–576 m (Cairns 1991; Cortés 2012).

***Polymyces wellsi* Cairns, 1991**

Fig. 74E

Material examined. AD4923: Co2951 (fragment).

Localities. Parrita Seep (~ 1037–1108 m).

Distribution. Originally described from the Galápagos Islands, 391–813 m (Cairns 1991), and thought to be cosmopolitan at depths of 355–1682 m (Cairns 2006).

Remarks. A new record for Costa Rica.

Cnidaria | Anthozoa | Hexacorallia | Scleractinia | Astrangiidae

***Astrangia* gen. inc.**

Fig. 74F

Material examined. AD4921: Co2943 (fragment).

Localities. Quepos Slide (~ 345–394 m).

Remarks. Damaged specimens, most likely *Astrangia*.

Cnidaria | Anthozoa | Octocorallia

We list entries following the phylogenetic relationships of Malacalcyonacea, Scleralcyonacea, and Pennatuloidae as defined in McFadden et al. (2022), and then alphabetically within those clades.

Cnidaria | Anthozoa | Octocorallia | Malacalcyonacea | Aquaumbriidae

***Aquaumbra klapferi* Breedy, van Ofwegen & Vargas, 2012**

Fig. 74G, H

Reference. Breedy et al. 2012**.

Additional material examined. S0219: MZUCR 3072 (voucher), Co3087 (tissue).

Localities. Rio Bongo Scar (610 m; this study).

Distribution. Originally described from the insular shelf of Isla del Coco, 268–308 m (Breedy et al. 2012) and subsequently reported from the West Florida Slope, Gulf of Mexico, 421 m (supplementary table 2 in Quattrini et al. (2020); Sequence Read Archive: SRS5647140, BioSample: SRS5647140).

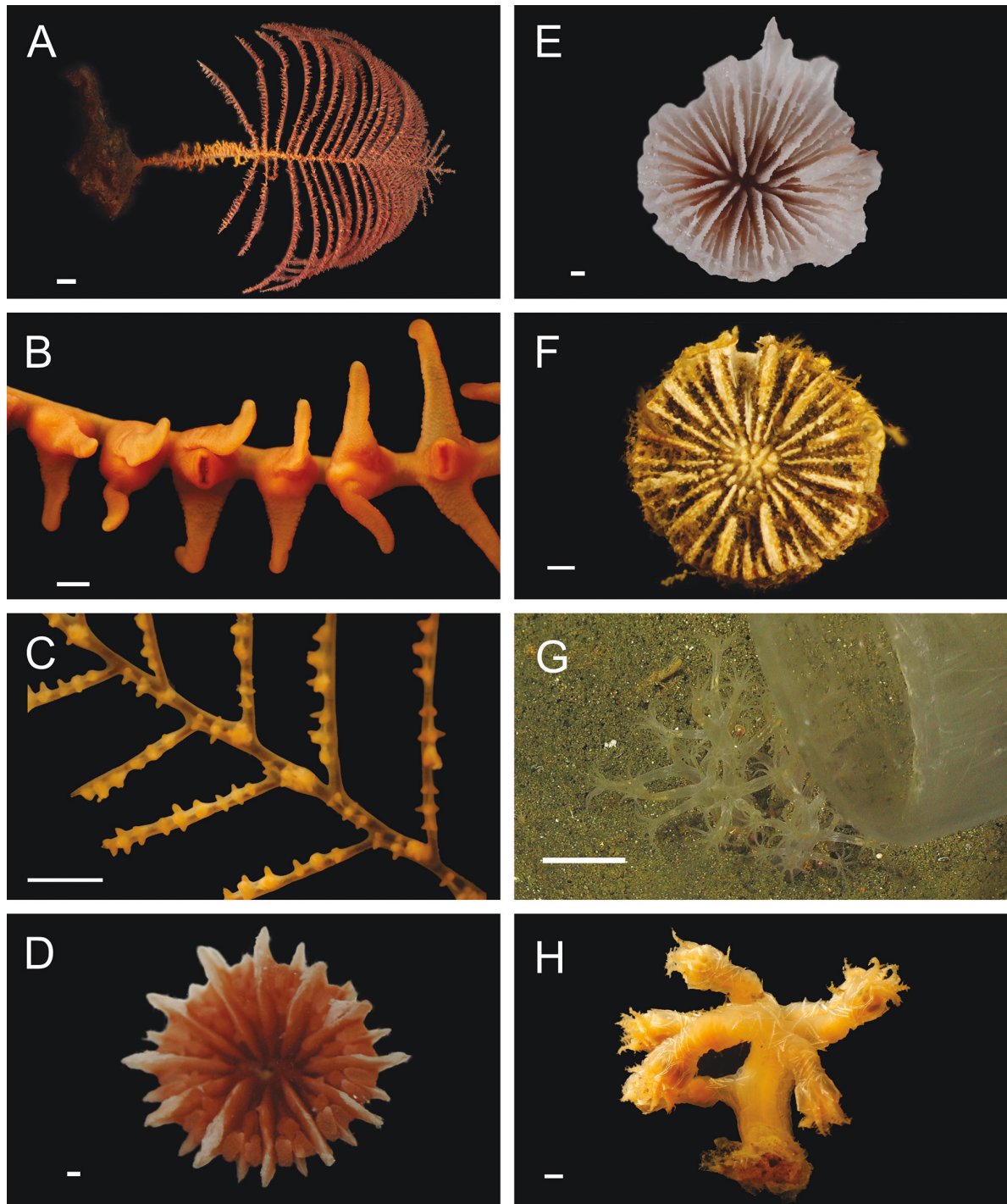


Figure 74. Cnidaria: Antipatharia, Scleractinia, and Octocorallia, representative live images **A** *Bathypathes stet.* (MZUCR 3541, colony) **B** *Bathypathes stet.* (MZUCR 3541, polyps) **C** *Lillipathes ritamariae* (Co2944) **D** *Javania cailleti* (Co2276) **E** *Polymyces wellsi* (Co2951) **F** *Astrangia* gen. inc. (Co2943) **G** *Aquaumbra klapferi* (MZUCR 3072, *in situ*). Credit: ROV SuBastian/Schmidt Ocean Institute **H** *Aquaumbra klapferi* (MZUCR 3072, detail). Scale bars: 1 cm (**A**, **G**); 1 mm (**B–F**, **H**).

The Gulf of Mexico record suggests that the relatively shallow eastern Pacific and western Atlantic populations were separated by the rise of the Isthmus of Panama relatively recently (see O’Dea et al. (2016)).

New records. The CRM specimen represents a new depth record for this species and establishes Rio Bongo Scar, 465 km northeast of Isla del Coco, as a second known locality in Costa Rica.

Cnidaria | Anthozoa | Octocorallia | Malacalcyonacea | Clavulariidae

Clavulariidae sp. MZUCR_3551

Fig. 75A, B

Material examined. AD4918: MZUCR 3551 (voucher), Co2929 (tissue).

Localities. Quepos Slide (338 m).

Remarks. The base of the colony was encrusted with hydroids and scallops, *Delectopecten vancouverensis* (M16184).

Clavulariidae sp. MZUCR_3552

Fig. 75C

Material examined. AD4923: MZUCR 3552 (voucher), Co2940 (tissue).

Localities. Parrita Seep (1075 m).

Cnidaria | Anthozoa | Octocorallia | Malacalcyonacea | Isididae

Isididae sp. MZUCR_3129

Fig. 75D, E

Material examined. AD4923: MZUCR 3129 (voucher), tissue samples Co2952, Co2954; Co2953.

Localities. Parrita Seep (1037 m).

Remarks. Host of *Heteroptychus galapagos* (C12816, C12817).

Isididae sp. MZUCR_3547

Fig. 75F

Material examined. AD4914: MZUCR 3547 (voucher), Co2927 (tissue).

Localities. Jacó Scar (1632 m).

Cnidaria | Anthozoa | Octocorallia | Malacalcyonacea | Paramuriceidae

***Acanthogorgia* sp. MZUCR_3113**

Fig. 75G, H

Material examined. AD4924: MZUCR 3113 (voucher), Co2956 (tissue).

Localities. Parrita Seep (1403 m).

Remarks. One colony was attached to a tube of *Lamellibrachia barhami*.

***Acanthogorgia* sp. MZUCR_3549**

Fig. 76A, B

Material examined. S0216: MZUCR 3549 (voucher), Co3079 (tissue).

Localities. Quepos Slide (316 m).

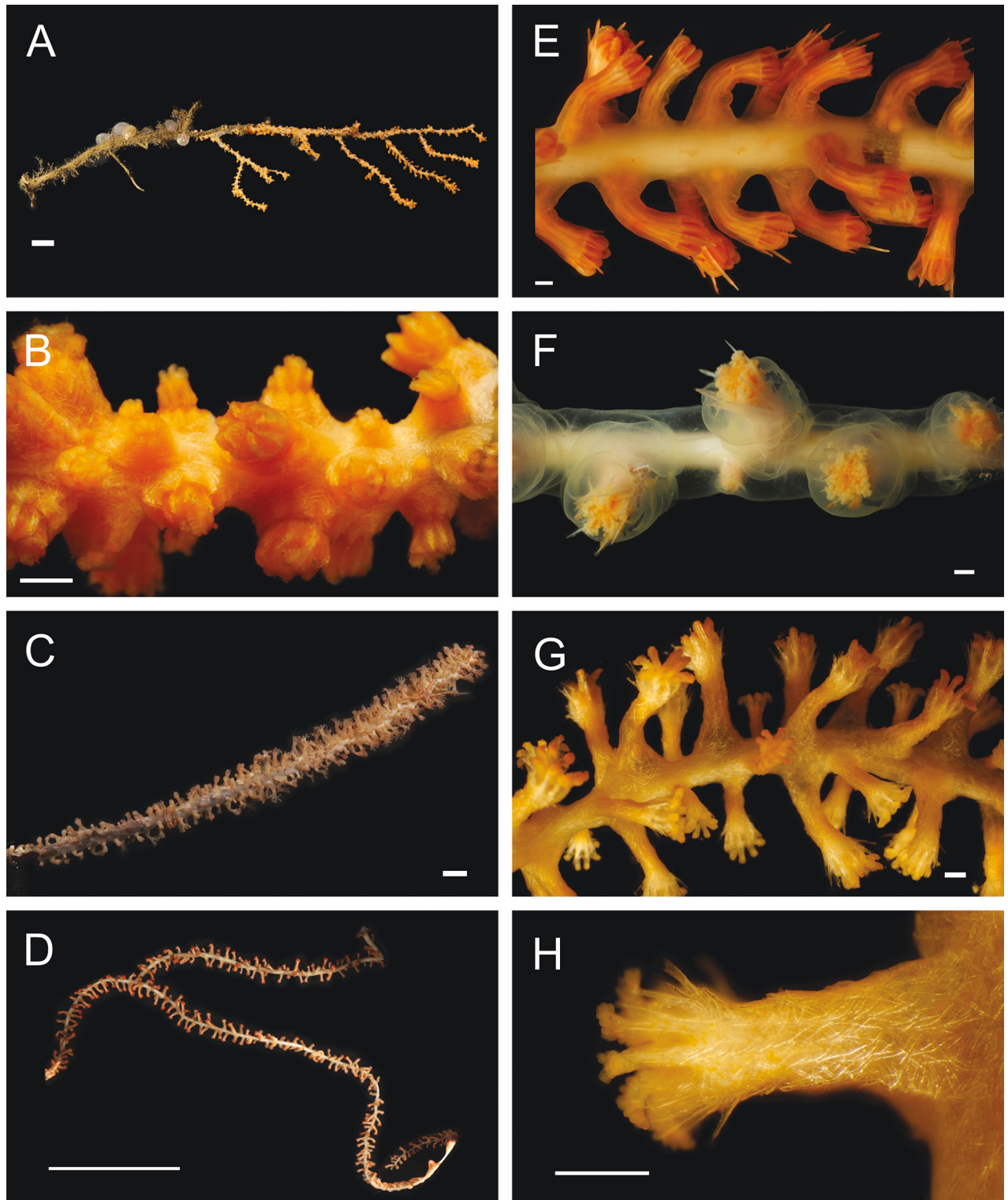


Figure 75. Cnidaria: Octocorallia, representative live images **A** Clavulariidae sp. MZUCR_3551 (MZUCR 3551, colony) **B** Clavulariidae sp. MZUCR_3551 (MZUCR 3551, polyps) **C** Clavulariidae sp. MZUCR_3552 (MZUCR 3552) **D** Isididae sp. MZUCR_3129 (MZUCR 3129, colony) **E** Isididae sp. MZUCR_3129 (Co2953, polyps) **F** Isididae sp. MZUCR_3547 (MZUCR 3547) **G** *Acanthogorgia* sp. MZUCR_3113 (MZUCR 3113, colony) **H** *Acanthogorgia* sp. MZUCR_3113 (MZUCR 3113, polyps). Scale bars: 1 cm (**A**, **C**); 1 mm (**B**, **E–H**); 10 cm (**D**).

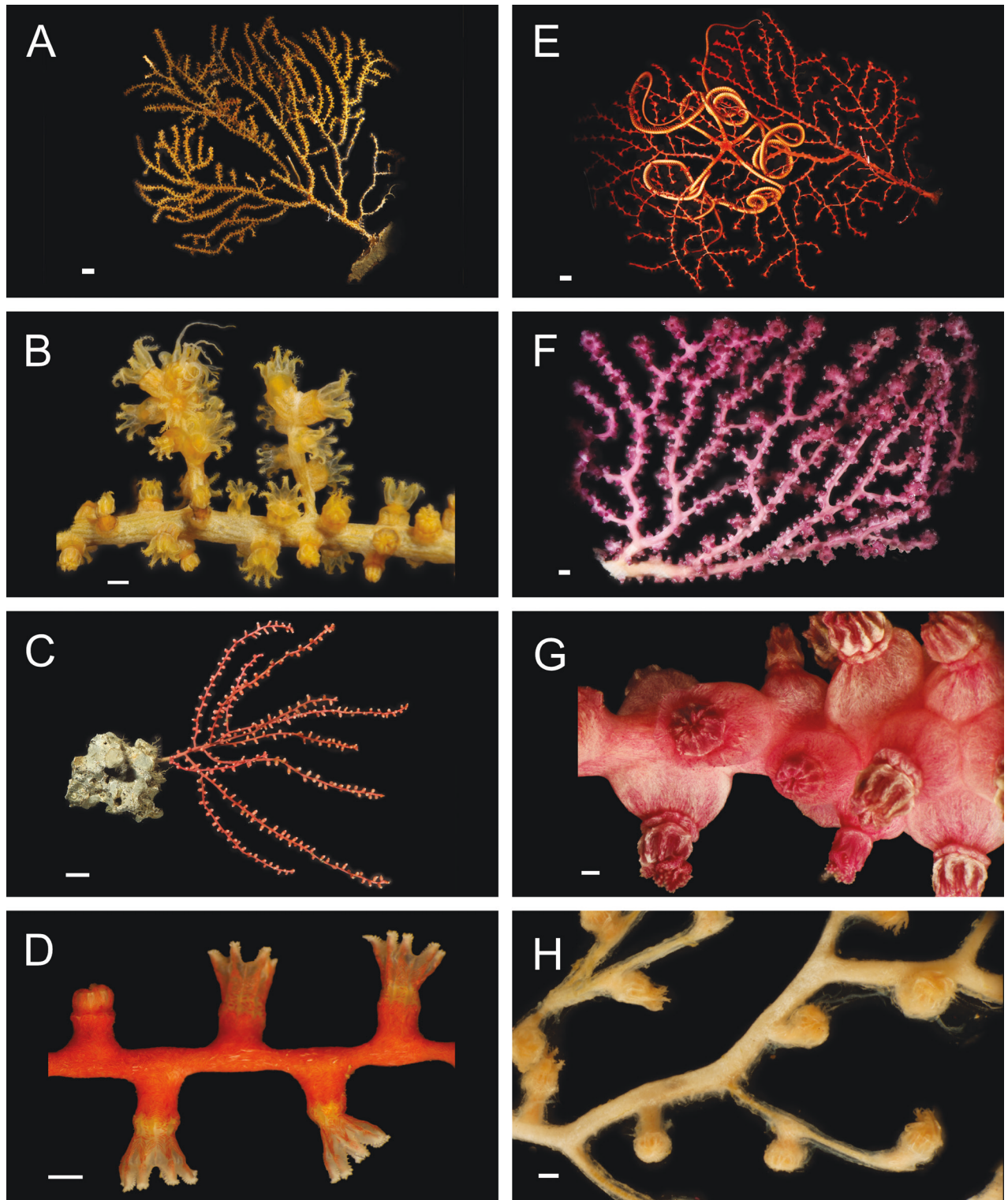


Figure 76. Cnidaria: Octocorallia, representative live images **A** *Acanthogorgia* sp. MZUCR_3549 (MZUCR 3549, colony) **B** *Acanthogorgia* sp. MZUCR_3549 (MZUCR 3549, polyps) **C** *Swiftia sahlingi* (MZUCR 2648, colony) **D** *Swiftia sahlingi* (Co2945, polyps) **E** *Swiftia stet.* (Co3080) **F** *Victorgorgia stet.* (MZUCR 3114, colony) **G** *Victorgorgia stet.* (MZUCR 3114, polyps) **H** *Chrysogorgia* sp. MZUCR_3057 (MZUCR 3057). Scale bars: 1 cm (**A**, **C**, **E**, **F**); 1 mm (**B**, **D**, **G**, **H**).

Cnidaria | Anthozoa | Octocorallia | Malacalcyonacea | Plexauridae

***Swiftia sahlingi* Breedy et al., 2019**

Fig. 76C, D

Reference. Breedy et al. 2019**.

Localities. Mound 12 (996–1002 m; type locality).

Distribution. Known only from the CRM seeps.

Remarks. Found in dense clusters on authigenic carbonates, often near sites of active methane seepage (Breedy et al. 2019).

***Swiftia* stet.**

Fig. 76E

Material examined. S0218: Co3080 (tissue).

Localities. Parrita Scar (1431 m).

Remarks. Associated with an ophiuroid, *Ophiocreas carnosus* (E7318). Possibly an undescribed species of *Swiftia*, distinct from *S. sahlingi*. Further taxonomic work on eastern Pacific *Swiftia* species is needed, including detailed comparison of this specimen to those from seamounts west of the CRM seeps at 2089–2270 m depth (Breedy et al. 2019).

Cnidaria | Anthozoa | Octocorallia | Malacalcyonacea | Victorgorgiidae

***Victorgorgia* stet.**

Fig. 76F, G

Material examined. AD4923: MZUCR 3114 (voucher), Co2939 (tissue).

Localities. Parrita Seep (1074 m).

Cnidaria | Anthozoa | Octocorallia | Scleralcyonacea | Chrysogorgiidae

***Chrysogorgia* sp. MZUCR_3057**

Fig. 76H

Material examined. AD4923: MZUCR 3057 (voucher), Co2948 (tissue).

Localities. Parrita Seep (1041 m).

***Chrysogorgia* sp. MZUCR_3063**

Fig. 77A

Material examined. AD4505: Co2270; AD4923: MZUCR 3063 (voucher), Co2950 (tissue).

Localities. Mound 11 (~ 1019–1025 m), Parrita Seep (1091 m).

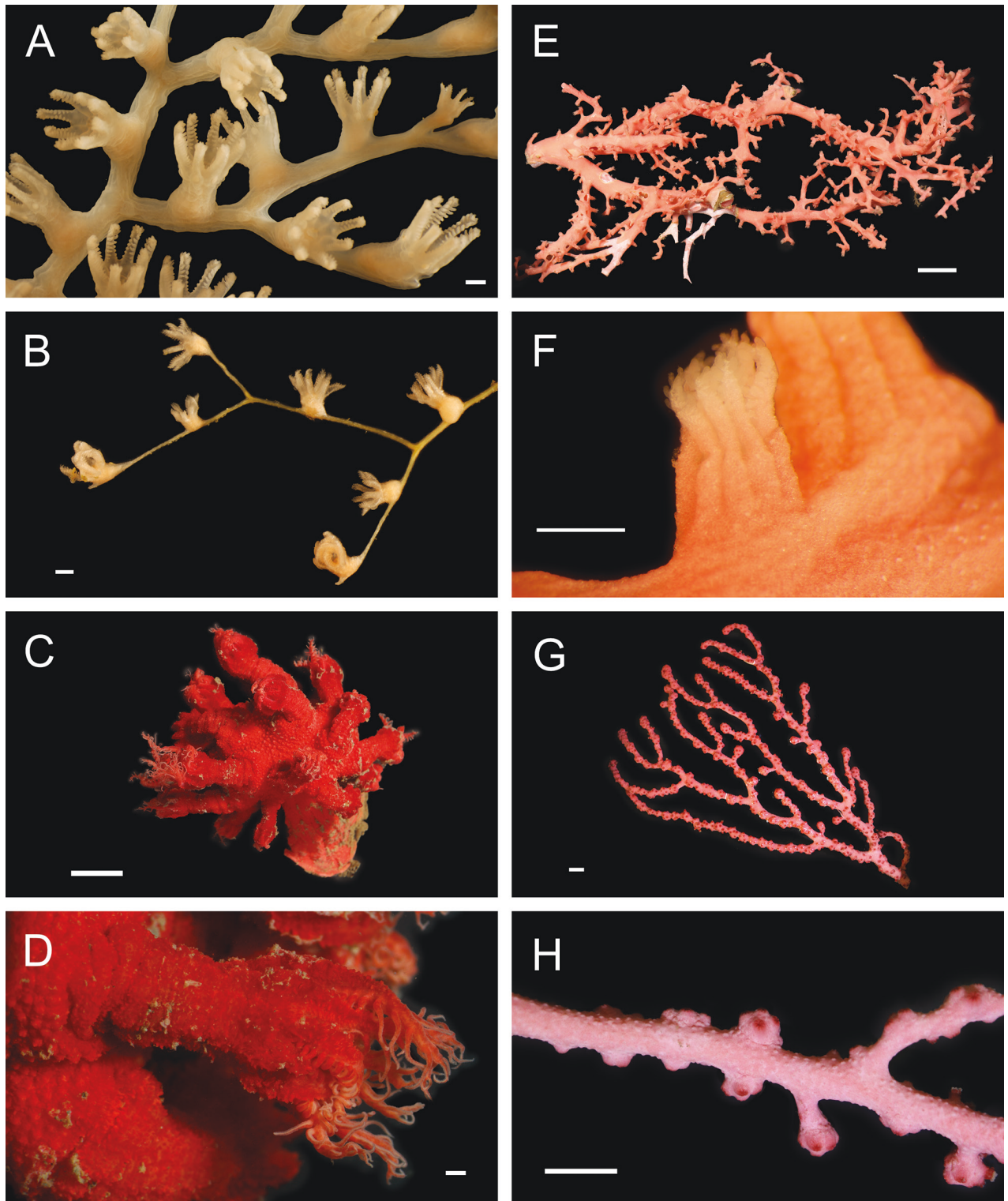


Figure 77. Cnidaria: Octocorallia, representative live images **A** *Chrysogorgia* sp. MZUCR_3063 (MZUCR 3063) **B** *Chrysogorgia* sp. MZUCR_3142 (MZUCR 3142) **C** *Anthomastus* stet. (Co2919, colony) **D** *Anthomastus* stet. (Co2919, polyps) **E** *Coralliidae* stet. (MZUCR 3130, colony) **F** *Coralliidae* stet. (MZUCR 3130, polyps) **G** *Paragorgia* stet. (Co3054, colony) **H** *Paragorgia* stet. (Co3054, detail). Scale bars: 1 mm (A, B, D, F); 1 cm (C, E, G, H).

***Chrysogorgia* sp. MZUCR_3142**

Fig. 77B

Material examined. AD4923: MZUCR 3142 (voucher), Co2949 (tissue).

Localities. Parrita Seep (~ 1037–1108 m).

Cnidaria | Anthozoa | Octocorallia | Scleralcyonacea | Coralliidae

***Anthomastus* stet.**

Fig. 77C, D

Material examined. AD4913: Co2919; MZUCR 3126 (voucher), Co2923 (tissue).

Localities. Jacó Scar (~ 1849–1867 m).

Coralliidae stet.

Fig. 77E, F

Material examined. AD4506: MZUCR 3550 (voucher), Co2271 (tissue); AD4923: MZUCR 3130 (voucher), Co2947 (tissue).

Localities. Parrita Seep (~ 1030–1094 m).

Remarks. Host of scaleworms, *Gorgoniapolyne* cf. *caeciliae* (A1549 on Co2271, A8455 on Co2947); a pyrgomatid barnacle (C12815 on Co2947); ophiuroids (*Ophiacantha moniliformis*, E4390 on Co2271; *Ophiuroglypha* cf. *meridionalis*, E7978 on Co2271); and anemones (Co2273 on Co2271, Co2874 on Co2947).

***Paragorgia* stet.**

Fig. 77G, H

Material examined. AD4990: Co3054 (tissue).

Localities. Parrita Seep (1407 m).

Remarks. Host of the nemertean *Alvinonemertes dariae* (Sagorny et al. 2022).

Cnidaria | Anthozoa | Octocorallia | Scleralcyonacea | Primnoidae

***Callogorgia* stet.**

Fig. 78A

Material examined. AD4914: MZUCR 3548 (voucher), Co2926 (tissue); AD4916: MZUCR 3539 (voucher), Co2931 (tissue); AD4923: MZUCR 3140 (voucher), Co2941 (tissue).

Localities. Parrita Seep (1052 m), Jacó Scar (~ 1604–1645 m).

Remarks. Co2941 was associated with an ophiuroid, *Ophiocreas carnosus* (E7072).

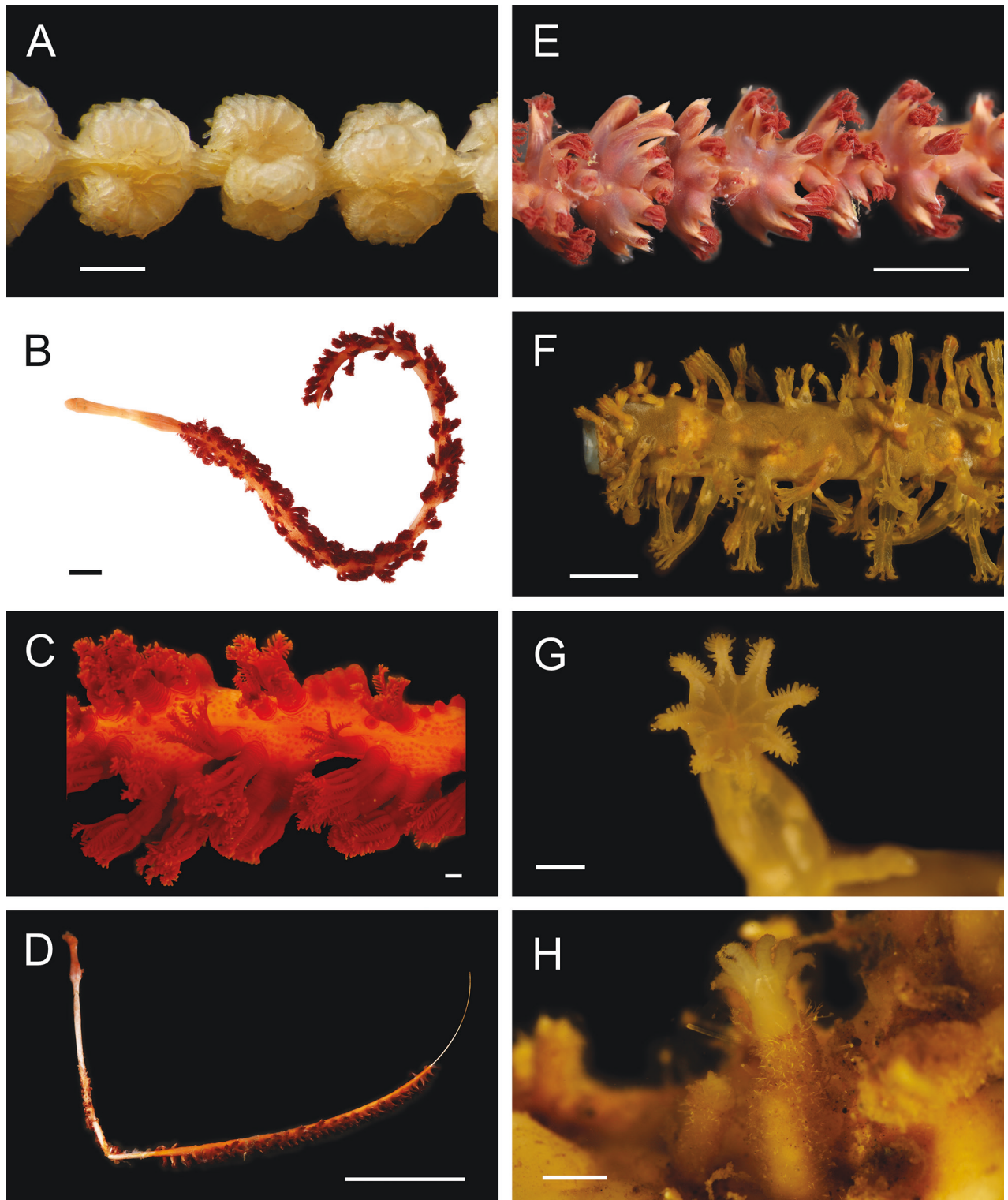


Figure 78. Cnidaria: Octocorallia, representative live images **A** *Callogorgia* stet. (MZUCR 3548) **B** *Anthoptilum grandiflorum* sp. inc. (MZUCR 3554, colony) **C** *Anthoptilum grandiflorum* sp. inc. (MZUCR 3554, polyps) **D** *Balticina* gen. inc. (MZUCR 3124, colony) **E** *Balticina* gen. inc. (MZUCR 3124, polyps) **F** *Octocorallia* sp. MZUCR_3112 (MZUCR 3112, colony) **G** *Octocorallia* sp. MZUCR_3112 (MZUCR 3112, polyps) **H** *Octocorallia* sp. MZUCR_3553 (MZUCR 3553). Scale bars: 1 mm (**A**, **C**, **G**, **H**); 1 cm (**B**, **E**, **F**); 10 cm (**D**).

Cnidaria | Anthozoa | Octocorallia | Scleralcyonacea | Pennatuloidae

***Anthoptilum grandiflorum* (Verrill, 1879) sp. inc.**

Fig. 78B, C

Material examined. AD4922: MZUCR 3554 (voucher), Co2937 (tissue).

Localities. Mound 12 (1005 m).

Distribution. Originally described from the north Atlantic off Nova Scotia and New England (Goode 1884), *A. grandiflorum* is considered cosmopolitan (Williams 2011).

Remarks. Associated with two squat lobsters, *Munidopsis hendersoniana* (C12809). Further morphological examination is needed to confirm the identification.

***Balticina* gen. inc.**

Fig. 78D, E

Material examined. AD4506: MZUCR 3538 (voucher), Co2274 (tissue); AD4923: MZUCR 3124.

Localities. Parrita Seep (1030–1098 m).

Cnidaria | Anthozoa | Octocorallia

***Octocorallia* sp. MZUCR_3112**

Fig. 78F, G

Material examined. AD4917: MZUCR 3112 (voucher), Co2934 (tissue).

Localities. Mound 12 (965 m).

Remarks. Stoloniferan morphology.

***Octocorallia* sp. MZUCR_3553**

Fig. 78H

Material examined. S0217: MZUCR 3553 (voucher), Co3083 (tissue).

Localities. The Thumb (~ 940–1070 m).

Remarks. Attached to a serpulid tube. Stoloniferan morphology.

***Octocorallia* sp. SIO_BIC_Co2278**

Fig. 79A, B

Material examined. AD4512: Co2278.

Localities. Quepos Slide (344 m).

Cnidaria | Hydrozoa | Anthoathecata

***Anthoathecata* stet.**

Fig. 79C

Material examined. AD4974: Co3016; S0215: Co3082.

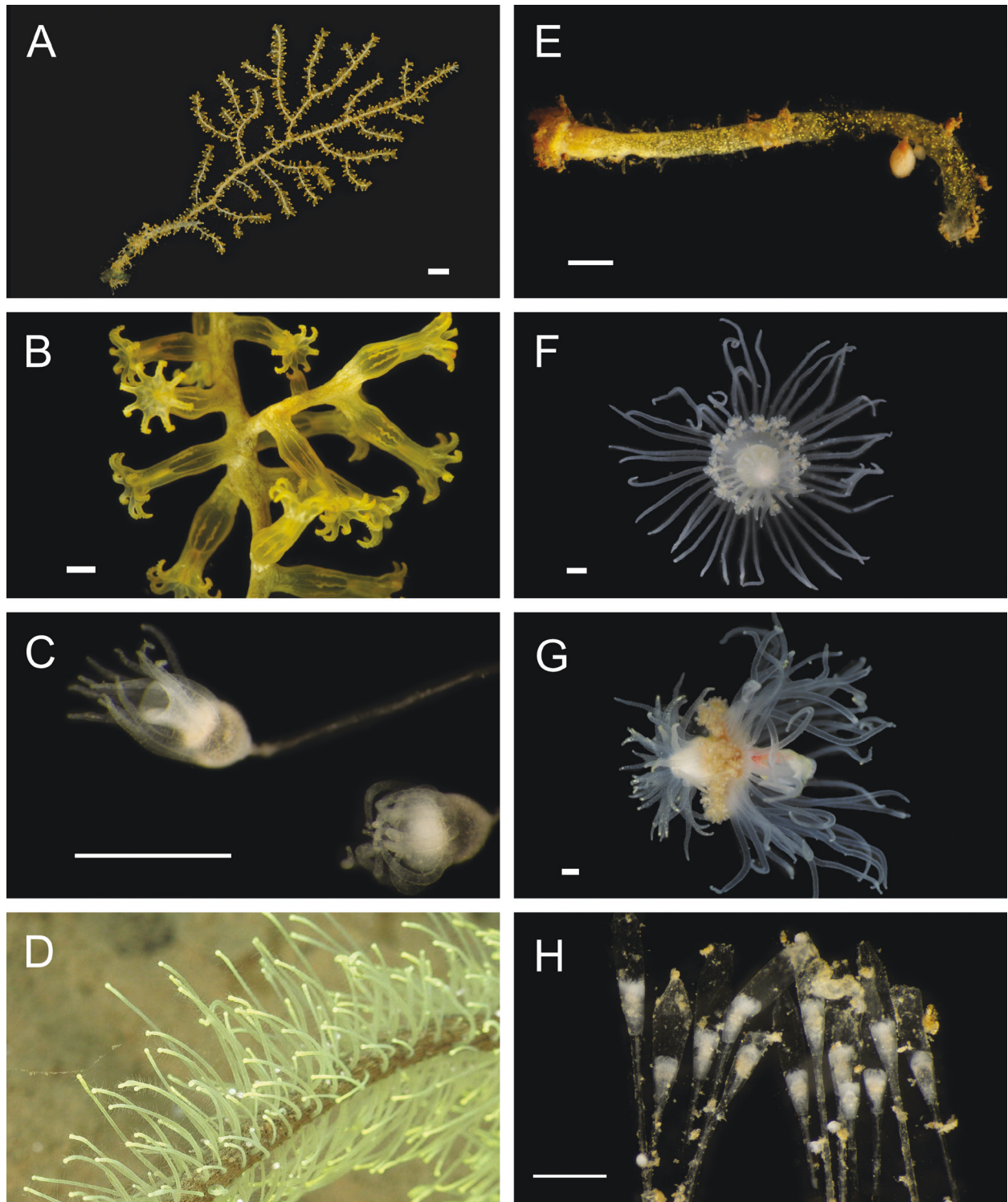


Figure 79. Cnidaria: Octocorallia and Hydrozoa, representative live images **A** Octocorallia sp. SIO_BIC_Co2278 (Co2278, colony) **B** Octocorallia sp. SIO_BIC_Co2278 (Co2278, polyps) **C** Anthothecata sp. (Co3016) **D** Candelabrum sp. (Co3088, *in situ* on the tube of *Melinnopsis* cf. *armipotens*, A12604). Credit: ROV SuBastian/Schmidt Ocean Institute **E** Candelabrum sp. (Co3088, polyp) **F** Corymorpha sp. (Co2272, oral view) **G** Corymorpha sp. (Co2272, lateral view) **H** Leptothecata sp. SIO_BIC_Co2928 (Co2928). Scale bars: 1 cm (**A**); 1 mm (**B–H**).

Localities. Mound 12 (1001–1011 m).

Remarks. Co3016 was associated with an experimentally deployed carbonate. Co3082 was associated with the tube of *Lamellibrachia donwalshi*.

Cnidaria | Hydrozoa | Anthoathecata | Aplanulata | Candelabridae

***Candelabrum* stet.**

Fig. 79D, E

Material examined. S0220: Co3088.

Localities. Subduction Plume (3503 m).

Remarks. Abundant on the exterior of a polychaete tube (*Melinnopsis* cf. *armipotens*, A12604), which extended above the seafloor to an estimated length of 20 cm. We thank Dhugal Lindsay (Japan Agency for Marine-Earth Science and Technology) for this identification.

Cnidaria | Hydrozoa | Anthoathecata | Aplanulata | Corymorphidae

***Corymorpha* stet.**

Fig. 79F, G

Material examined. AD4506: Co2272.

Localities. Parrita Seep (~ 1030–1179 m).

Remarks. We thank Dhugal Lindsay for this identification.

Cnidaria | Hydrozoa | Leptothecata

***Leptothecata* sp. SIO_BIC_Co2928**

Fig. 79H

Material examined. AD4916: Co2928.

Localities. Jacó Scar (1854 m).

Remarks. Epibionts on the fecampiid egg case Pt64.

Cnidaria | Scyphozoa | Coronamedusae | Coronatae | Nausithoidae

***Nausithoe* stet.**

Fig. 80A

Material examined. AD4918: Co2972.

Localities. Quepos Slide (338 m).

Remarks. The outer surfaces of the tubes of these polyps show prominent rings with sharp edges and relatively sparse sculpturing, indicative of *Nausithoe* type 1 in Jarms et al. (2002). Differentiation of *Nausithoe* species based on polyp morphology alone is difficult (Molinari et al. 2023). The tube-dwelling polyps of coronate scyphozoans (Atorellidae and Nausithoidae) have been reported from near-surface waters to at least 3,233 m (Song et al. 2021; Molinari et al. 2023). We thank Xikun Song (Institute of Deep-Sea Science and Engineering, Chinese Academy of Sciences) and Dhugal Lindsay for this identification.

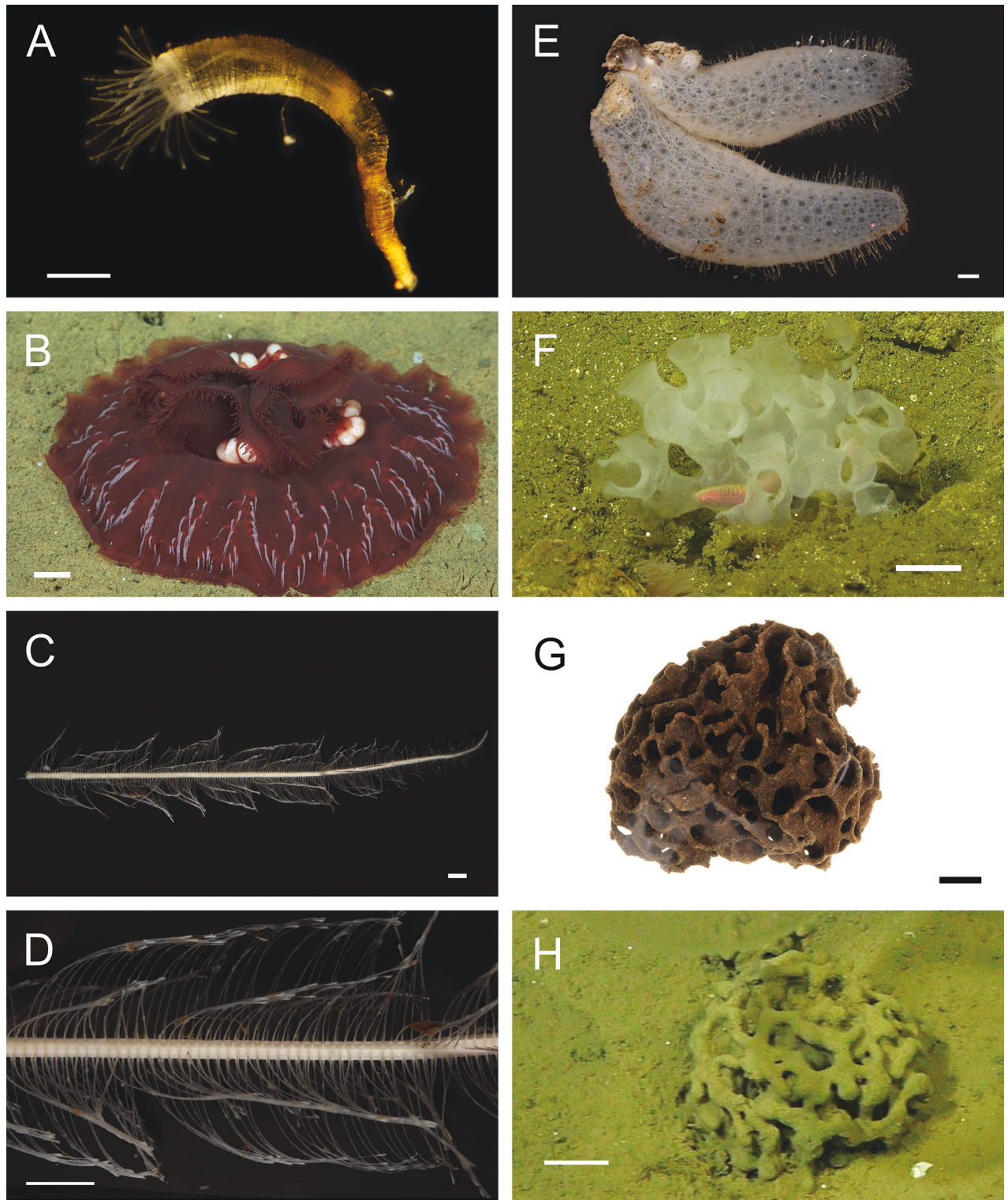


Figure 80. Cnidaria: Scyphozoa, Porifera, and Foraminifera, representative live images **A** *Nausithoe* stet. (Co2972, polyp) **B** *Ulmaridae* stet. (Co3011, medusa *in situ*). Credit: ROV SuBastian/Schmidt Ocean Institute **C** *Asbestopluma* gen. inc. (P1754) **D** *Asbestopluma* gen. inc. (P1754, detail) **E** *Dictyaulus* gen. inc. (P1687) **F** *Farrea occa* (P1753, *in situ* with polynoid scaleworms A10096 and A10098 inside). Credit: ROV SuBastian/Schmidt Ocean Institute **G** *Reticulammina* stet (B11172) **H** *Shinkaiya* stet. (*in situ*; specimen destroyed). Credit: ROV SuBastian/Schmidt Ocean Institute. Scale bars: 1 mm (**A**); 1 cm (**B–H**).

Cnidaria | Scyphozoa | Discomedusae | Semaestomeae

Ulmaridae *stat.*

Fig. 80B

Material examined. S0220: Co3011 ([PQ300674](#)) (tissue; voucher disintegrated during collection).

Localities. Subduction Plume (3434 m).

Remarks. This specimen was observed alive yet upside down on the seafloor. We thank George Matsumoto (Monterey Bay Aquarium Research Institute) and Rebecca Helm (Georgetown University) for generating the COI sequence. This medusa showed morphological and perhaps behavioral similarities to an undescribed hadal ulmarid that was reported from 8200 m in the New Britain Trench and observed skimming the sediment surface to feed on particulates (Gallo et al. 2015) (Dhugal Lindsay, pers. comm. 25 November 2019).

Porifera

We thank Lonny Lundsten (Monterey Bay Aquarium Research Institute) for the identification of these specimens.

Porifera | Demospongiae | Poecilosclerida | Cladorhizidae

***Asbestopluma* gen. inc.**

Fig. 80C, D

Material examined. S0220: P1754 (tissue).

Localities. Subduction Plume (3601 m).

Remarks. Associated with a scaleworm, *Macellicephala* sp. SIO_BIC_A10099 (A10099).

Porifera | Hexactinellida | Lyssacinosa | Euplectellidae

***Dictyaulus* gen. inc.**

Fig. 80E

Material examined. AD4923: P1687.

Localities. Parrita Seep (1098 m).

Remarks. Host of stenopodids, *Spongioloides galapagensis* (C12735).

Porifera | Hexactinellida | Sclerophora | Farreidae

***Farrea occa* Bowerbank, 1862**

Fig. 80F

Material examined. S0219: P1752, P1753.

Localities. Rio Bongo Scar (602–631 m).

Distribution. Considered cosmopolitan at depths of 204–1901 m, with several subspecies occurring in the eastern Pacific between California and Panama, 523–1244 m (Lopes et al. 2011).

Remarks. Host of polynoid scaleworms (A10096, A10098).

Chromista

Chromista | Foraminifera | Monothalamea | Xenophyophoroidea

Reticulammina stet.

Fig. 80G

Reference. Levin and Rouse 2020.

Additional material examined. AD4913: BI1172.

Localities. Jacó Scar (1866–1885 m).

Remarks. One specimen (destroyed) was associated with snailfish eggs (*Paraliparis* stet., BI1369), representing one of the first two recorded uses of xenophyophores as nursery habitat by fishes (Levin and Rouse 2020).

Shinkaiya stet.

Fig. 80H

Reference. Levin and Rouse 2020.

Localities. Mound Jaguar (1902 m).

Remarks. This specimen (destroyed) was associated with snailfish eggs (*Acantholiparis* stet., BI1371) (Levin and Rouse 2020).

Syringammina stet.

Fig. 81A

Material examined. AD4913: BI1158.

Localities. Jacó Scar (1885 m).

Chromista | Foraminifera

Foraminifera stet.

Fig. 81B

Material examined. AD4921: BI1153, BI1164; S0219: BI1366.

Localities. Quepos Slide (~ 345–394 m), Rio Bongo Scar (661 m).

Remarks. Arborescent foraminiferans associated with naturally occurring wood falls.

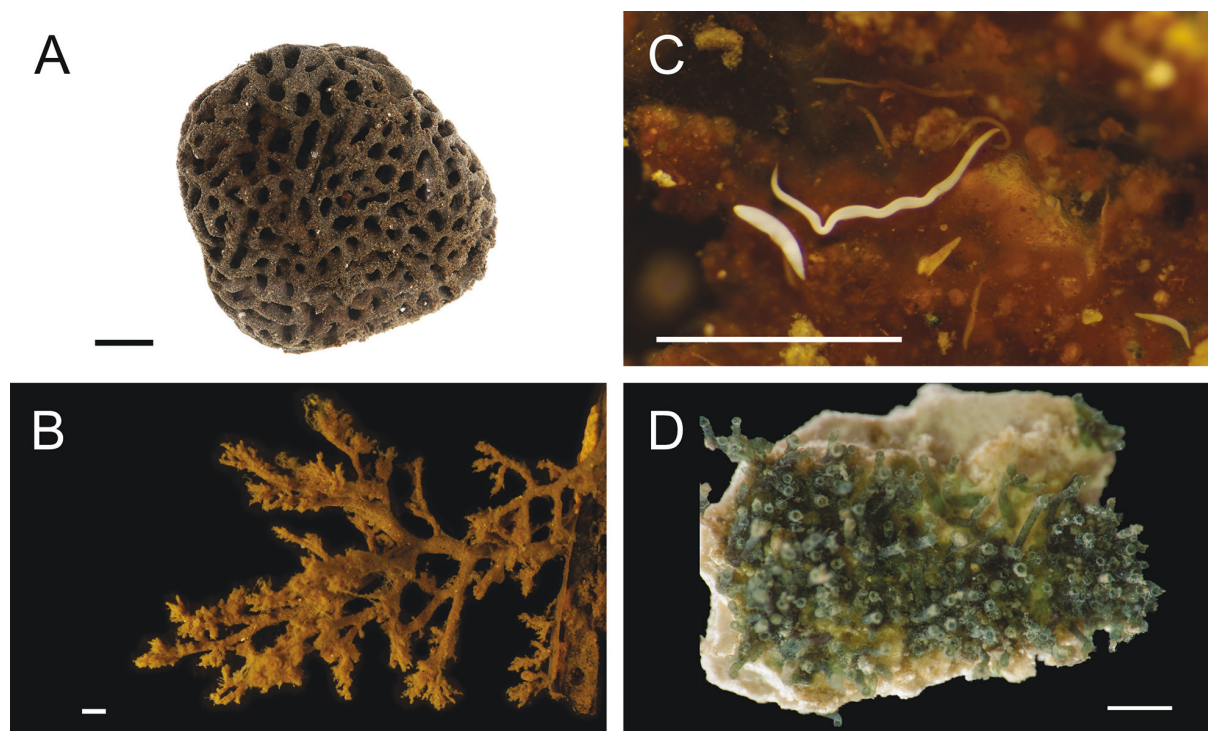


Figure 81. Foraminifera and Ciliophora, representative live images **A** *Syringammina stet.* (BI1158) **B** Foraminifera stet. (BI1164) **C** Ciliophora stet. (BI1173) **D** *Eufolliculina caerulea* (BI1051). Scale bars: 1 cm (**A**); 1 mm (**B–D**).

Ciliophora

Ciliophora stet.

Fig. 81C

Material examined. AD4918: BI1173.

Localities. Quepos Slide (~ 333–408 m).

Remarks. Motile ciliates were associated with a bacterial film on a carbonate rock from a site of active seepage.

Eufolliculina caerulea Pasulka & Rouse in Pasulka et al. 2017

Fig. 81D

Reference. Pasulka et al. 2017**.

Localities. Mound 12 (~ 1000 m).

Distribution. Also known from eastern Pacific seeps off southern California (type locality: Split Ridge, off San Diego, 865 m) and the Guaymas Basin, total depth range 865–1565 m (Pasulka et al. 2017).

Results

Taxonomic identification

We report 488 species-level taxa (i.e., distinct taxonomic units in the sense of morphospecies, though often distinguished by genetics rather than morphology)

occurring at the CRM seeps, including 131 described (formally named) species and at least 58 undescribed species new to science (Table 3, Suppl. material 1: table S1). The remaining 299 species-level taxa present some degree of uncertainty in their identification. The 488 total taxa include 155 annelids, 96 crustaceans, 42 cnidarians, 55 echinoderms, 108 mollusks, 9 nemerteans, 17 other animals, and 6 chromists. Of the 131 described species, 49 were originally described from the CRM seeps.

New records

We report 15 range extensions for species occurring in the eastern Pacific: five new northern records, nine new southern records, and one new eastern Pacific record (Table 4, Suppl. material 1: table S1). We also report 16 new depth records (6 new minimum depths, 10 new maximum depths), and 3 new seep records of species known to occur at vents or organic falls (Table 4, Suppl. material 1: table S1).

Biogeographic ranges

Of the 131 described species occurring at the CRM seeps, 38 are known only from Costa Rica/Panama, 64 occur elsewhere in the eastern Pacific, six also occur through the northern Pacific to Japan, six occur more broadly in the Pacific, and 17 occur in multiple ocean basins (Table 5, Suppl. material 1: table S1).

Occurrences at other chemosynthesis-based habitats

Of the 131 described species occurring at the CRM seeps, 13 also occur at hydrothermal vents, 14 also occur at organic falls, and six occur at all three types of chemosynthesis-based habitats (Table 6, Suppl. material 1: table S1).

Table 3. Summary of taxa occurring at the CRM seeps. “Undescribed species” indicates species that have been identified with certainty as new to science but have not yet been formally described and named. “Taxa with taxonomic uncertainty” indicates distinct taxonomic units that could not be identified with certainty to the level of species, e.g., “*Parougia* cf. *sulleyi*” or “*Chaetognatha* stet.”

Higher taxonomy	# described species (# which were originally described from the CRM seeps)	# undescribed species noted in this work	# taxa with taxonomic uncertainty	Total	# species with new biogeographic records at the CRM seeps
Annelida	38 (27)	35	82	155	2
Arthropoda	30 (4)	4	62	96	4
Cnidaria	5 (2)	0	37	42	0
Echinodermata	20 (1)	9	26	55	5
Mollusca	28 (7)	8	72	108	4
Nemertea	6 (6)	1	2	9	0
Other Animalia	3 (1)	1	13	17	0
Chromista	1 (1)	0	5	6	0
Total	131 (49)	58	299	488	15

Table 4. New biogeographic records. New northern and southern records are indicated by the location of the previously reported limit. New depth records were defined as a minimum difference of 100 m from a previously reported minimum or maximum depth. New seep records pertain to species previously reported from vents or organic falls. Only occurrences reported in peer-reviewed publications were included in these assessments; other occurrences of potential interest, e.g., unpublished data or GBIF records, are discussed in the text.

Higher taxonomy	Species	Known biogeographic range (including this work)	New biogeographic records reported in this work	Known depth range (including this work)	New minimum depth record	New maximum depth record	New seep records reported in this work
Annelida	<i>Lindaspio dibranchiata</i>	Gulf of California to CRM	new southern record (previously Guaymas Basin only)	820–2008 m	820 m (previously 2004 m)		new seep record (previously from sedimented vents)
Annelida	<i>Myzostoma josefinae</i>	central California to CRM	new southern record (previously to the Guaymas Basin)	1020–1878 m		1878 m (previously ~ 1350 m)	
Arthropoda	<i>Alvinocaris costaricensis</i>	CRM only		742–1800 m	742 m (previously ~ 1000 m)		
Arthropoda	<i>Heteroptychus galapagos</i>	CRM to Galápagos	new northern record (previously Galápagos only)	1012–1037 m			
Arthropoda	<i>Lebbeus scrippsi</i>	Gulf of California to northern Chile		768–1768 m		1768 m (previously 1245 m)	
Arthropoda	<i>Munidopsis alvisca</i>	Juan de Fuca Ridge to CRM	new southern record (previously to the Guaymas Basin)	1401–2008 m	1401 m (previously 1545 m)		new seep record (previously from vents)
Arthropoda	<i>Spongicoloides galapagensis</i>	CRM to Galápagos	new northern record (previously Galápagos only)	717–1091 m		1091 m (previously 717 m)	
Arthropoda	<i>Sternostylus defensus</i>	CRM to Galápagos	new northern record (previously Galápagos only)	717–1210 m			
Cnidaria	<i>Aquaumbra klapferi</i>	CRM to Isla del Coco; Gulf of Mexico		268–610 m		610 m (previously 421 m)	
Echinodermata	<i>Araeosoma leptaleum</i>	California to Galápagos		740–1271 m		1271 m (previously 1106 m)	
Echinodermata	<i>Calamocrinus diomedae</i>	CRM to Galápagos		717–1659 m		1659 m (previously 1430 m)	
Echinodermata	<i>Ophiacantha moniliformis</i>	southern California to CRM	new southern record (previously to southern Mexico)	377–1244 m			
Echinodermata	<i>Ophiocreas carnosus</i>	CRM to southern Chile	new northern record (previously central Chile)	175–1632 m		1632 m (previously 983 m)	
Echinodermata	<i>Ophioleuce gracilis</i>	Pacific, widely distributed		1710–2882 m	1710 m (previously ~ 2000 m)		
Echinodermata	<i>Ophiomitra partita</i>	central Mexico to CRM	new southern record (previously Islas Mariás only)	1236–1887 m		1887 m (previously 1236 m)	
Echinodermata	<i>Ophiophrura liodisca</i>	western Pacific; CRM	new eastern Pacific record (previously Japan to New Zealand)	606–1620 m			
Echinodermata	<i>Thrissacanthias penicillatus</i>	Washington to CRM	new southern record (previously to the Gulf of California)	55–1503 m			
Mollusca	<i>Belknapchiton halistreptus</i>	southern Mexico to CRM	new southern record (previously southern Mexico only)	609–3436 m	609 m (previously 902 m)		

Higher taxonomy	Species	Known biogeographic range (including this work)	New biogeographic records reported in this work	Known depth range (including this work)	New minimum depth record	New maximum depth record	New seep records reported in this work
Mollusca	<i>Delectopecten vancouverensis</i>	Alaska to CRM; Bering Sea to Sea of Japan	new southern record (previously to Baja California)	20–4100 m			
Mollusca	<i>Ennucula colombiana</i>	CRM to Peru		11–997 m		997 m (previously 734 m)	
Mollusca	<i>Eulimella lomana</i>	southern California to CRM	new southern record (previously to the Guaymas Basin)	1168–2008 m			
Mollusca	<i>Tindariopsis grasslei</i>	Gulf of California to CRM		992–2012 m	992 m (previously 1400 m)		
Mollusca	<i>Tripoplax balaenophila</i>	CRM to central Chile	new northern record (previously central Chile only)	240–344 m		344 m (previously 240 m)	new seep record (previously from bone)
		Total	15		6	10	3

Table 5. Known biogeographic ranges of the described species occurring at the CRM seeps. Numbers indicate the count of described species within each category.

Eastern Pacific		Northern range limit			
		Costa Rica/ Panama	Mexico	California	Juan de Fuca region
Southern range limit	Costa Rica/Panama	38	10	10	7
	Ecuador	7	2	1	0
	Peru/northern Chile	3	8	3	2
	Central Chile	4	1	1	1
	Southern Chile	1	1	1	1
Northern Pacific (Japan to Costa Rica/Panama)		6			
Pacific, broadly		6			
Multiple ocean basins		17			

Table 6. Known occurrences at other chemosynthesis-based habitats for described species occurring at CRM seeps. The count of species in each category is indicated in the bottom row. New seep records of species previously reported from vents or organic falls are indicated in bold. The new seep records were collected from naturally occurring seep substrates rather than wood or bone deployments.

Seeps + vents	Seeps + vents + organic falls	Seeps + organic falls
<i>Amphisamytha fauchaldi</i>	<i>Archivesica gigas</i>	<i>Alvinonemertes christianeae</i>
<i>Archinome levinae</i>	<i>Escarpia spicata</i>	<i>Escondidacntrainea panamensis</i>
<i>Calyptogena diagonalis</i>	<i>Eulimella lomana</i>	<i>Heterocarpus vicarius</i>
<i>Bathymodiolus thermophilus</i>	<i>Phreagena soyoae</i>	<i>Micospina auribohorum</i>
<i>Calyptogena costaricana</i>	<i>Provanna laevis</i>	<i>Munidopsis carinipes</i>
<i>Galapagomystides veranae</i>	<i>Xyloplax princealberti</i>	<i>Munidopsis hendersoniana</i>
<i>Lamellibrachia barhami</i>		<i>Myzostoma josefinae</i>
<i>Lepetodrilus guaymasensis</i>		<i>Osedax frankpressi</i>
<i>Lindaspio dibranchiata</i>		<i>Osedax knutei</i>
<i>Munidopsis alvisca</i>		<i>Peinaleopolynoe elvisi</i>
<i>Phreagena extenta</i>		<i>Peinaleopolynoe mineoi</i>
<i>Provanna ios</i>		<i>Provanna pacifica</i>
<i>Shinkai longipedata</i>		<i>Tripoplax balaenophila</i>
		<i>Typhlonida propinqua</i>
13	6	14

Discussion

Biodiversity at the CRM seeps

The CRM seeps support a biodiverse fauna and have been a rich site of species discovery. Of the 131 described species occurring at these seeps, 49 were newly described from this area (Table 3, Suppl. material 1: table S1), beginning with the vesicomid clam *Calyptogena diagonalis* in 1999 (Barry and Kochevar 1999). Of these 49 species, 47 (excluding *C. diagonalis* and *C. costaricana*) were originally described from the cruises summarized in this work. Many of the 58 undescribed species identified in this study will be formally described in due course. With further work, it may be possible to identify the remaining 299 “species” reported with taxonomic qualifiers as known matches to described species, or to ascertain that they are additional undescribed species.

The extensive field sampling program in this study has been essential for these discoveries. For the 488 taxa reported to occur at the CRM seeps, all occurrence records included material collected during the five cruises of this study, except for five previously published vesicomid clam species: *Archivesica* sp. 7 sec. Audzijonyte et al. 2012, *Calyptogena costaricana*, *C.* sp. 3 sec. Audzijonyte et al. 2012, *C.* sp. mt-V sec. Goffredi et al. 2003, and *Phreagena extenta*. These five species were collected during prior work at deeper seep sites northwest of those targeted in this study (Peek et al. 2000; Goffredi et al. 2003; Audzijonyte et al. 2012). Further biological sampling at the northwestern seeps and others at comparable depths (3000–4000 m; e.g., Subduction Plume) may elucidate whether these clams are restricted by depth and/or local habitat conditions.

Comparison to checklists for Costa Rica

An extensive catalog has been compiled for the overall marine biodiversity of Costa Rica by Wehrtmann and Cortés (2009b). At least 4690 marine species are known from the entire Pacific coast, with most records from shallow waters (Cortés 2012). This study adds six species to the checklist of 75 echinoderms reported from Costa Rican waters at > 200 m depth (Alvarado et al. 2022) (*Ophiacantha inconspicua*, *Ophiacantha moniliformis*, *Ophiocreas carnosus*, *Ophiomitra partita*, *Ophiophrura liodisca*, *Xyloplax princealberti*). This study also adds 21 species to the 139 crustaceans recorded from the Costa Rican Pacific at > 200 m depth (Azofeifa-Solano and Cortés 2020) (*Colossendeis macerrima*, *Eucopia sculpticauda*, *Heteroptychus galapagos*, *Lebbeus scrippsi*, *Lycaea pulex*, *Mesopleustes abyssorum*, *Munidopsis agassizii*, *Munidopsis alvisca*, *Munidopsis carinipes*, *Munidopsis cortesi*, *Munidopsis girguisi*, *Munidopsis granosicorium*, *Munidopsis hendersoniana*, *Munidopsis similis*, *Nematocarcinus faxoni*, *Oncopagurus haigae*, *Paracrangon areolata*, *Parapagurus foraminosus*, *Spongicoloides galapagensis*, *Sternostylus defensus*, *Typhlonida propinqua*). Previous deep-sea work based on the 2009 and 2010 cruises (Levin et al. 2012, 2015) included lists of taxa in the supplemental material. We have referenced those records where identifications could be confidently matched to extant voucher material; many of those listings reflect provisional identifications based on morphology and have since been revised to reflect taxonomic work on reference specimens at SIO-BIC.

Comparison to global seep checklists

The 131 described species recorded from the CRM seeps represent a substantial portion of the ~ 600 species recorded from all > 100 seep sites worldwide as of 2011 (German et al. 2011), and of 863 species currently recorded from all chemosynthetic ecosystems worldwide (Ramírez-Llodra 2024). With increased discovery and biological characterization of seeps in recent decades, this global list has expanded approximately threefold. As of 1998, only 211 species (of which 116 were named) had been reported from all 24 seep sites known worldwide at the time (Sibuet and Olu 1998).

Comparison to seep-adjacent background habitats

Mobile fauna known from non-seep “background” habitats can utilize and aggregate at seeps (Carney 1994; Sibuet and Olu 1998; Tunnicliffe et al. 1998; Niemann et al. 2013; Levin et al. 2016). As discussed by Carney (1994), any attempt to categorize fauna as “vagranths, colonists, or endemics” with respect to seeps depends on an accurate and thorough inventory of the surrounding environment, which falls outside the scope of this study. We therefore did not assign such categorizations to the fauna reported in this study, and we echo caution in interpreting fauna as seep-endemic or regionally endemic based on absence of evidence.

Several species reported from the shallower CRM seeps in this work (Quepos Seep, Rio Bongo Scar, and Jacó Summit) were previously well known from shelf and slope depths (< 350 m) inshore of the seeps, e.g., the commercially harvested shrimp *Heterocarpus vicarius* and abundant bycatch crustaceans: *Grimothea monodon*, *Plesionika trispinus*, and *Squilla biformis* (Wehrtmann and Echeverría-Sáenz 2007; Wehrtmann et al. 2010). Studies of seeps on the northern California slope (~ 500 m depth) (Levin et al. 2000) and the Koryak slope in the Bering Sea (400–700 m) (Rybakova et al. 2022) have also reported strong similarity between seep and non-seep faunas, remarking that the seep fauna at these depths may be pre-adapted to the organic-rich reducing conditions at seeps (Levin et al. 2000).

We acknowledge the importance of future work on the non-seep habitats of the CRM, especially seamounts (Wehrtmann and Cortés 2009a; Cortés 2012). Several seamounts occur at similar depths as the deeper seeps, e.g., Quepos Plateau and Seamount 1 at 1410–2184 m (Hatch et al. 2020; Sagorny et al. 2022), and share geological characteristics with seeps such as Jacó Scar, Parrita Scar, and Rio Bongo Scar (Sahling et al. 2008). The CRM seep sediments also show spatial gradients of macrofaunal assemblages and geochemistry, described in detail in Ashford et al. (2021b). Sampling in this work was centered on sites of active seepage and their transition zone (tens to hundreds of meters) into the surrounding environment, but more extensive sampling of far-transition and background sites (~ 1 km and beyond) at various localities would help to evaluate the full extent of this transition zone.

Comparison to other seep, vent, and organic fall sites

Although site-specific taxonomic inventories are not available for many seeps and direct comparisons are complicated by differences of methodology and scope, the count of 488 distinct morphospecies reported at the CRM seeps

in this study exceeds those published for other seep sites to date. At least 335 macrofaunal morphospecies occur at the Koryak slope seeps in the Bering Sea (Rybakova et al. 2022), 289 at the Laptev Sea seeps in the Siberian Arctic (Vedenin et al. 2020), 134 at Hydrate Ridge off Oregon (Levin et al. 2017), 101 at the Concepción seeps off central Chile (Sellanes et al. 2008), 83 at the seeps of El Pilar off Trinidad and Tobago (Amon et al. 2017), 65 at the Haima seeps in the South China Sea (He et al. 2023), 65 at the Gulf of Mexico seeps (Bergquist et al. 2003), 60 at the Del Mar seep off southern California (Grupe et al. 2015), 60 at the Taitao Peninsula seep off southern Chile (Zapata-Hernández et al. 2014), 50–60 at the Florida Escarpment and Blake Ridge seeps in the western Atlantic (Turnipseed et al. 2003), 30 at the Guaymas Basin seeps (Portail et al. 2015), and 20 at the South Chamorro Seamount serpentinite-hosted seep on the Mariana Forearc (Chen et al. 2024c).

Recent macrofaunal inventories from vents have reported 130 morphospecies from the Piip Volcano in the Bering Sea (Rybakova et al. 2023), 92 from the Galápagos Rift (Chen et al. 2024b), 91 from the Okinawa Trough vent system (Tunnicliffe et al. 2024), 12–70 at each of ten other western Pacific vent systems (Tunnicliffe et al. 2024), 62 from the Loki's Castle Vent Field on the Arctic Mid-Ocean Ridge (Eilertsen et al. 2024), and 29 from the Amami Rift vent field off southern Japan (Chen et al. 2024a). Individual whale falls off California have been reported to support 36–190 macrofaunal species (Baco and Smith 2003; Lundsten et al. 2010). Experimental wood deployments in the Gulf of Mexico (McClain et al. 2023) and southwestern Atlantic (Saeedi et al. 2019) have been reported to support 44 and 114 macrofaunal invertebrate species, respectively.

Depth ranges at the CRM seeps

At the CRM seeps, pairs of morphologically indistinguishable yet genetically distinct species can be separated by depth. The serpulid tubeworms *Laminatebus joycebrooksae* and *L. paulbrooksi* occur at ~ 1000 m and ~ 1400–2400 m, respectively (Rouse and Kupriyanova 2021). The chemosymbiont-bearing mussels *Bathymodiolus nancyschneiderae* (1000–1100 m) and *B. billschneideri* (1400–1900 m) occupy non-overlapping depths, yet the morphologically distinct *B. earlougheri* co-occurs with both species across their combined depth range (McCowan et al. 2020). Furthermore, the four species of symbiotic scaleworms (*Branchiopolynoe* spp.) that occupy the mantle cavity of these mussels show no apparent depth or host specificity (Lindgren et al. 2019). Although the effects of depth are difficult to disentangle from those of seep geochemistry and other parameters, these examples show that a putative morphospecies can comprise at least two depth-segregated cryptic species, requiring careful taxonomy and potentially different management considerations.

Range extensions reinforce eastern Pacific biogeographic connections

The range extensions in this study reinforce the biogeographic connections of the CRM seeps to neighboring regions of the eastern Pacific, from the Juan de Fuca Ridge to southern Chile. The smallest range extensions were new northern records of species previously known only from the Galápagos

(representing ~ 8 degrees of latitude and ~ 1,100 km). The greatest eastern Pacific range extension was the new northern record of *Tripoplax balaenophila*, previously known only from central Chile (representing ~ 45 degrees of latitude and ~ 5,100 km).

Mexico and Gulf of California

The Middle America Trench connects the CRM to seeps off western Mexico. The East Pacific Rise (EPR) links the Cocos Plate to seeps and vents in the Gulf of California (Kimura et al. 1997; Fiedler and Lavín 2006), including those of the Guaymas Basin (Portail et al. 2015) and the Pescadero Transform Fault (Goffredi et al. 2017). Of the 131 described species at the CRM, 79 species (60%) also range to Mexico or beyond, and 10 of these species are presently known only from the CRM and Mexico. The nine new southern records reported in this work represent range extensions from Mexican waters, and we expect that further deep-sea sampling in both regions will reveal additional overlap. In particular, the Guaymas Basin seeps at ~ 2000 m share several high-biomass, habitat-forming species with the CRM seeps (vestimentiferan tubeworms *Lamellibrachia barhami* and *Escarpia spicata*, vesicomyid clams *Archivesica gigas* and *Phreagena soyoeae*) (Portail et al. 2015), suggesting broad similarity of the two communities.

Galápagos and Coco Submarine Volcanic Range

The Galápagos seamounts, rising from the abyssal plain at ~ 3000 m, are connected to Costa Rica by the Coco Submarine Volcanic Range, which was formed by the Galápagos Hotspot and subducts beneath the CRM (Kimura et al. 1997; von Huene et al. 2000; Fiedler and Lavín 2006). The seamount subduction section of the CRM shares the characteristic geochemistry of the Galápagos seamounts (von Huene et al. 2000; Sahling et al. 2008), suggesting biogeographical affinities as well. Of the 131 described species at the CRM, 67 species (51%) also range to the Galápagos region or beyond (including five northern range extensions), and seven species are presently known only from the CRM and Galápagos.

The first systematic ROV characterization of deep-sea benthic invertebrate communities at the Galápagos seamounts (290–3373 m) reported 70 morphospecies based on collection of 90 specimens and extensive imagery (Salinas-de-León et al. 2020). Thirty of those morphospecies were undescribed and new to science at the time, including the squat lobster *Heteroptychus galapagos*, which we report here from the CRM seeps as a range extension. These examples illustrate the extent of undiscovered biodiversity along the Coco Submarine Volcanic Range and the importance of FAIR taxonomic inventories (Howell et al. 2020) in describing this biodiversity. Under circumstances where certain data cannot be shared publicly, e.g., permit conditions noted in Salinas-de-León et al. (2020), datasets from biogeographically connected regions such as the CRM will be important for studies of comparative biodiversity.

Located approximately halfway between the Galápagos seamounts and the CRM, Isla del Coco (Cocos Island) is thought to have high rates of endemism, with 1688 marine species recorded and 747 of these not found elsewhere in

Pacific Costa Rica (Cortés 2012). The deepest regions of Isla del Coco (> 450 m) remain under-explored as they are beyond the capability of the locally operating DeepSee submersible and may reveal further biogeographic linkages along the Coco Submarine Volcanic Range. For example, the soft coral *Aquaumbra klapferi* was previously known from Isla del Coco, but here we also report it from Rio Bongo Scar, establishing a new depth record and an association with seeps.

Peru-Chile margin

Seeps supporting diverse fauna occur along the Peru-Chile margin (Olu et al. 1996; Sellanes et al. 2008), and this region is an important focus area for further biological investigation (Ramírez Llodra et al. 2003; German et al. 2011; Miloslavich et al. 2011; Tapia-Guerra et al. 2021; Wagner et al. 2021). Of the 131 described species at the CRM, 57 species (44%) also range to Peru or beyond, including two northern range extensions. Additional taxonomic inventories along the Peru-Chile margin (e.g., Valentich-Scott et al. 2020) will be instrumental in addressing biogeographic knowledge gaps, such as the extent of connectivity between the eastern Pacific and Antarctica.

Trans-Pacific and broader ranges

Six of the 131 described species in this work (5%) reportedly also range to Japan, e.g., via the Alaska margin. Four of these species have been confirmed to show trans-Pacific genetic connectivity (*Archivesica gigas*, *Phreagena extenta*, *Phreagena soyoae*, *Provanna laevis*), whereas range-wide genetic data are not currently available for the others (*Delectopecten vancouverensis*, *Shinkai longipedata*).

Twenty-three of the described species in this work (18%) reportedly have even wider distributions, extending into the southwestern Pacific or other ocean basins. Some of these examples are corroborated by genetics (e.g., *Osedax frankpressi*, *Munidopsis similis*), whereas others are suspected to represent cryptic species complexes or synonymies warranting resurrection (e.g., *Ophiura flagellata*).

Other biogeographic connections

An important geological factor in the evolutionary history of the CRM seeps is the separation of the Atlantic and Pacific Oceans by the rise of the Isthmus of Panama (O’Dea et al. 2016). For example, the polychaete *Amphisamytha fauchaldi* from the CRM seeps is most closely related to the Atlantic *A. lutzii* (Desbruyères & Laubier, 1996) rather than to any other Pacific *Amphisamytha* (Stiller et al. 2013). Similarly, *Lamellibrachia donwalshi* from the CRM seeps is more closely related to a clade of Atlantic species, including *L. anaximandri* Southward, Andersen & Hourdez, 2011, than to any Pacific *Lamellibrachia* (McCowan and Rouse 2018). The soft coral *Aquaumbra klapferi* has been reported from relatively shallow eastern Pacific and western Atlantic waters (Breedy et al. 2012; Quattrini et al. 2020), consistent with these populations being separated by the rise of the Isthmus of Panama relatively recently compared to deeper-water species.

Where several congeners occur at the CRM seeps, the CRM species are not necessarily sister taxa to one another, and their true relationships reveal a variety of biogeographic affinities. For example, *Lamellibrachia barhami* is more closely related to *L. satsuma* Miura, 1997 from the western Pacific than to *L. donwalshi* (McCowin and Rouse 2018). *Bathymodiolus nancyschneiderae* is more closely related to a western Pacific *Bathymodiolus* clade than to *B. billschneideri* and *B. earlougheri*, which are sister taxa and form an eastern Pacific clade with *B. thermophilus* (also occurring at the CRM seeps) and *B. antarcticus* (McCowin et al. 2020). As with depth ranges, the patterns for the four CRM seep *Branchiopolynoe* species do not track those of their mussel hosts: *B. halliseyae* and *B. kajsae* are sister taxa and most closely related to *B. seepensis* from the western Atlantic, whereas *B. eliseae* and *B. meridae* belong to an eastern Pacific clade along with *B. symmytilida* from the EPR (Lindgren et al. 2019). Together, these examples indicate evolutionary linkages from the CRM to the EPR, western Atlantic, and western Pacific.

Intersection of seeps, vents, and organic remains

The first submersible-based accounts of deep-sea fauna at seeps (Paull et al. 1984) and whale remains (Smith et al. 1989) immediately noted the similarity to hydrothermal vent communities. Subsequent comparisons of these three ecosystems have depended on comprehensive lists of well identified taxa (Sibuet and Olu 1998; Tunnicliffe et al. 2003; Watanabe et al. 2010; Bernardino et al. 2012; Kiel 2016).

The biogeographic and evolutionary connections among these ecosystems are complex. Hydrothermal vents and seeps can occur in close geographical proximity, as known from the Guaymas Basin (Simoneit et al. 1990; Portail et al. 2015), Edison Seamount off Papua New Guinea (Herzig et al. 1998), and several regions of Japan (Watanabe et al. 2010). Vents and seeps may intersect with organic falls where they converge with forested coasts, kelp beds, or whale migration routes (Smith and Baco 2003; Bernardino et al. 2012). Furthermore, sedimented vents such as those at Manus Basin, the Guaymas Basin, Escanaba Trough at Gorda Ridge, and the Middle Valley on the Juan de Fuca Ridge, are considered intermediate environments by combining the soft substrate and fluid chemistry characteristic of seeps with the elevated temperatures and metal concentrations characteristic of bare-rock vents (Tunnicliffe et al. 1998, 2003; Levin et al. 2009; Bernardino et al. 2012; Kiel 2016).

Overlap of species occurrences

Worldwide, at least ~ 700 morphospecies occur at hydrothermal vents and at least 407 occur at organic remains, compared to ~ 600 at seeps (Smith and Baco 2003; German et al. 2011; Chapman et al. 2019). Comparative studies attempting to control for these conditions and other variables have suggested that seep communities tend to show higher diversity than comparable vent communities (Turnipseed et al. 2003; Bernardino et al. 2012; Portail et al. 2015).

Globally, at least 24 species co-occur at seeps and vents (Portail et al. 2015), at least 20 co-occur at seeps and organic remains (Smith and Baco 2003), and at least 11 co-occur at organic remains and vents (mainly sedimented vents at

the Guaymas Basin and the Juan de Fuca Ridge) (Smith and Baco 2003). At least eight species occur at all three habitat types: *Calyptogena elongata*, *Calyptogena pacifica*, *Chiridota heheva*, *Cocculina craigsmithi*, *Eulimella lomana*, *Escarpia spicata*, *Idas washingtonia*, *Pyropelta corymba*, and *P. musaica* (Smith et al. 1989; Tunnicliffe et al. 1998; Smith and Baco 2003; Thomas et al. 2020). At least two of these eight species occur at the CRM seeps (*Eulimella lomana* and *Escarpia spicata*; *Pyropelta* cf. *corymba* and *P.* cf. *musaica* require genetic confirmation). In this study, we also add two species to the list of overlap between seeps and vents (*Lindaspio dibranchiata*, *Munidopsis alvisca*), and one species (*Tripoplax balaenophila*) to the list of overlap between seeps and whale remains (Table 6).

Modes of symbiosis

Modes of chemosynthetic symbiosis can be important for explaining and predicting the occurrence of symbiont-bearing taxa across chemosynthesis-based habitats. At the CRM seeps, all three species of vestimentiferan tubeworms host the same thiotrophic symbiont, whereas the three numerically dominant mussels each host a unique thiotrophic symbiont (Brzechffa and Goffredi 2021). Methanotrophic examples from the CRM seeps include the serpulid worm *Laminatubus paulbrooski* and the sabellid worm *Bispira* sp. SIO_BIC_A9587 (Goffredi et al. 2020; Rouse and Kupriyanova 2021). Farming of epibiotic bacteria (likely thiotrophic or methanotrophic) by the yeti crab *Kiwa puravida* represents another conspicuous mode of symbiosis at the CRM seeps (Thurber et al. 2011; Goffredi et al. 2014).

Areas for further work

Despite our intensive and collaborative sampling efforts, we acknowledge that this inventory is still surely incomplete.

Geochemistry

The seep sites in this study were located in the central CRM spanning the “mound segment” and the “seamount subduction segment” (Sahling et al. 2008), thereby covering habitats with different geochemical characteristics. Faunal sampling at additional seeps, especially at the distal ends of each segment, may reveal additional diversity and suggest new hypotheses about the ecology and biogeography of this region.

Depth

The depths of sampling locations in this study (~ 300–3600 m) span the known bathymetric range of the entire CRM (Sahling et al. 2008) and approximately half the known depth range for seeps globally (~ 300–7400 m) (Sibuet and Olu 1998; Fujikura et al. 1999; Watanabe et al. 2010). Most of the sampling effort in this study occurred between 400–1800 m (Table 1), so further sampling of the deepest seeps may yield additional new records, species, and ecological findings.

Pelagic linkages

This study focused on benthic habitats, but the influence of seep and vent plumes on benthopelagic and pelagic fauna warrants further investigation (Levin et al. 2016). The opportunistically collected pelagic samples in this study have provided vouchered reference sequences for animals that are generally challenging to collect (e.g., the holopelagic sea cucumber *Pelagothuria natatrix*). The few pelagic fauna encountered on the seafloor (pyrosomes and an undescribed scyphomedusa) offer potential examples of trophic linkage between the CRM seeps and the overlying water column.

Conclusions

Importance of regional taxonomic inventories

Regional taxonomic inventories provide essential baseline biodiversity data for ecosystem science, monitoring, and management. Species occurrence data form the foundation of trait databases and biogeographic delineations, all of which can inform the design of effective ecological reserves (Van Dover et al. 2012; Chapman et al. 2019). Consequently, knowledge gaps in taxonomy and biogeography are a major impediment to deep-sea conservation, particularly of invertebrates (Cardoso et al. 2011; Ramírez-Llodra et al. 2011; Glover et al. 2018; Sigwart et al. 2023).

This inventory adds to the extensive existing documentation of marine biodiversity in Costa Rica, including recent emphasis on the deep sea (Wehrtmann and Cortés 2009a; Alvarado et al. 2010, 2012, 2022; Cortés 2012; Azofeifa-Solano and Cortés 2020). Much of the rich terrestrial and marine biodiversity of Costa Rica, including 50% of the coastline and an area of seamounts surrounding Isla del Coco National Park, has received conservation status as Protected Areas or other designations (Alvarado et al. 2012).

Recent quantitative economic research has demonstrated that Costa Ricans place high value on their seeps, with a willingness to pay for seep protection being highest for programs that protect seeps with endemic species (Pereira et al. 2024). In a related survey of deep-sea care and understanding, Costa Ricans ranked the provision of habitat and biodiversity as their most important connection to the deep sea (Pereira et al. 2024). Thus, continued biodiversity discovery is significant to the people of Costa Rica and to the stewardship of their deep marine ecosystems. We hope that the biodiversity data in this work will inform future marine policy in Costa Rica and the tropical eastern Pacific.

With thousands of seep sites and hundreds of vent sites estimated to exist across the deep sea (Levin et al. 2016), the ability to make direct and repeatable biodiversity comparisons is important for global research and policy-building. We advocate for the inclusion of genetic data in taxonomic inventories to enhance the robustness of taxonomic identifications and to allow evaluation of species ranges and assessment of population connectivity. The combination of museum voucher specimens, morphologically informative images, and diagnostic DNA sequences establishes critical reference libraries for broader scientific,

educational, and policy uses (e.g., Glover et al. 2015). For example, the abyssal fauna of the Clarion-Clipperton Zone are being inventoried in a growing body of specimen-based, DNA-supported taxonomic work (Dahlgren et al. 2016; Glover et al. 2016; Christodoulou et al. 2020; Bribiesca-Contreras et al. 2022).

Recommendations

For taxonomic inventories to maximize their potential, we emphasize the importance of careful specimen processing and molecular taxonomy, supported by deep submergence technology and biological collections.

Submersibles enable high-quality genetic and morphological analyses through targeted collection, precise data logging, high-resolution *in situ* imaging, and live recovery of intact animals. These vehicles, equipped with appropriate sampling and imaging tools, are particularly important for work on deep chemosynthetic ecosystems which are not well suited to trawls and other non-selective sampling gear. Reliable access to, and support for, deep-sea technologies remain essential to the advancement of deep-sea research (Liang et al. 2021; Marlow et al. 2022; Miller and Virmani 2023).

During specimen processing at sea, clear documentation and judicious field protocols are critical for scientific and legal traceability (Gemeinholzer et al. 2010; Kroupa and Remsen 2010; Templado et al. 2010; Glover et al. 2015; Howell et al. 2020). For example, unique identifiers (alphanumeric codes, ideally pre-printed on indelible labels) are important for linking physical samples to metadata, digital sequence information, and permissible downstream applications. Not only are voucher specimens critical for the integrity of taxonomic and genomic research, but it is important to record the relationship of the voucher to the tissue used to generate sequence data (i.e., same individual or a proxy) (Pleijel et al. 2008; Buckner et al. 2021). These field practices facilitate the deposition of well-curated physical specimens and their associated data into accessible repositories.

Biological collections have been central to this study and serve as foundations for biodiversity research, among other scientific and societal benefits (Suarez and Tsutsui 2004; National Science and Technology Council Committee on Science Interagency Working Group on Scientific Collections 2009; National Academies of Sciences Engineering and Medicine 2020). We intend for this inventory to underscore the importance of biological collections and to serve as an integrative resource for future studies of biodiversity and biogeography at deep-sea ecosystems.

Acknowledgements

We are grateful to the captains and crew of R/V Atlantis and R/V Falkor and the pilots and technicians of HOV Alvin and ROV SuBastian for crucial operations at sea and collecting specimens. We thank the science parties of cruises AT15-44 (2009), AT15-59 (2010), AT37-13 (2017), AT42-03 (2018), and FK190106 (2019); members of the Rouse laboratory; collection assistants at SIO-BIC; and Rita Vargas and colleagues at MZUCR for their support processing specimens

and data. We appreciate artists-at-sea Lily Simonson and Rebecca Rutstein for their interdisciplinary interpretations of the Costa Rica seep ecosystems. Many taxonomic collaborators are thanked, in the applicable sections of this work, for specific identifications and advice. We thank Christopher Glasby, Nathalie Yonow, and an anonymous reviewer for thoughtful and constructive feedback on this manuscript.

Additional information

Conflict of interest

The authors have declared that no competing interests exist.

Ethical statement

No ethical statement was reported.

Funding

Field collection (2009–2018) and laboratory analyses were funded by U.S. National Science Foundation (NSF) grants: OCE-0826254 (Rouse, Levin), OCE-0939557 (Rouse, Levin), OCE-1634172 (Rouse, Levin), and OCE-1635219 (Cordes). Field collection (2019) and article processing charges were funded by the Schmidt Ocean Institute. We thank the Bohn family for support of DNA sequencing. We thank Scripps Institution of Oceanography Director Margaret Leinen and the Director's Office for continued support of SIO-BIC core operations.

Author contributions

GWR, HC, and CAS conceptualized the project. GWR designed the methodology. GWR, LAL, and EEC secured funding and ship time. EEC and LAL served as chief scientists for the cruises. EEC and JC secured collection permits. JC obtained authorization for the genetic components of this project and led the ongoing collaboration with the University of Costa Rica. EEC, JC, SKG, LAL, and GWR supervised field operations. Shipboard specimen identification was led by GWR with contributions from OSA, OB, EEC, JC, SKG, ASH, LAL, AKM, MFM, CAS, JS, ET, AW, and NGW. Live specimen photography, documentation, and preservation were led by GWR along with ASH, AKM, MFM, CAS, JS, ET, and NGW. Specimens were vouchered and catalogued by HC and CAS. For most specimens, JIC, HC, ASH, and MFM performed DNA extraction and sequencing, and GWR led the analysis of the sequences to arrive at identifications. Taxon-specific DNA extraction, sequencing, analysis, and identification were conducted by KMK (Aplacophora) and TDO (Ophiuroidea). Morphological identifications, biogeographic and taxonomic expertise, and/or additional phylogenetic analyses were contributed by: STA (Galatheaidea), OSA (Peracarida), OB (Anthozoa), JC (Anthozoa), DJE (Polyplacophora), MEH (Decapoda), NMK (Echinoidea), KMK (Aplacophora), CLM (Asteroidea), AKM (Holothuroidea), RM (Echinoidea), TDO (Ophiuroidea), FP (Phyllodoidea), GWR (Annelida), PVS (Bivalvia), JS (Terebelliformia), ET (Sabellidae), AW (Gastropoda), MKW (Decapoda), and NGW (Gastropoda). Haplotype networks were prepared by CAS and GWR. CAS wrote the initial draft, prepared figures, and coordinated contributions from all authors with input from GWR. All authors reviewed and approved the final version.

Author ORCIDs

Charlotte A. Seid  <https://orcid.org/0000-0002-5307-691X>
Avery S. Hiley  <https://orcid.org/0000-0001-8956-5080>
Marina F. McCowin  <https://orcid.org/0000-0001-9674-5557>
José I. Carvajal  <https://orcid.org/0000-0003-2255-9705>
Shane T. Ah Yong  <https://orcid.org/0000-0002-2820-4158>
Oliver S. Ashford  <https://orcid.org/0000-0001-5473-7057>
Odalisca Breedy  <https://orcid.org/0000-0001-5686-4164>
Douglas J. Eernisse  <https://orcid.org/0000-0002-5457-7484>
Shana K. Goffredi  <https://orcid.org/0000-0002-9110-9591>
Michel E. Hendrickx  <https://orcid.org/0000-0001-9187-6080>
Kevin M. Kocot  <https://orcid.org/0000-0002-8673-2688>
Christopher L. Mah  <https://orcid.org/0000-0002-0178-8237>
Allison K. Miller  <https://orcid.org/0000-0002-5334-2771>
Nicolás Mongiardino Koch  <https://orcid.org/0000-0001-6317-5869>
Rich Mooi  <https://orcid.org/0000-0003-1533-8488>
Timothy D. O'Hara  <https://orcid.org/0000-0003-0885-6578>
Josefin Stiller  <https://orcid.org/0000-0001-6009-9581>
Ekin Tilic  <https://orcid.org/0000-0003-0463-322X>
Paul Valentich-Scott  <https://orcid.org/0000-0003-0019-7643>
Mary K. Wicksten  <https://orcid.org/0000-0002-9097-353X>
Nerida G. Wilson  <https://orcid.org/0000-0002-0784-0200>
Erik E. Cordes  <https://orcid.org/0000-0002-6989-2348>
Lisa A. Levin  <https://orcid.org/0000-0002-2858-8622>
Jorge Cortés  <https://orcid.org/0000-0001-7004-8649>
Greg W. Rouse  <https://orcid.org/0000-0001-9036-9263>

Data availability

Sequences are available on GenBank and linked to BioProject PRJNA1142170 (CRM specimens, 274 new sequences) or PRJNA1136884 (comparative material from other localities, 5 new sequences). SIO-BIC specimen records are available and continually updated through the online collection database (<https://sioapps.ucsd.edu/collections/bi/>). The SIO-BIC specimen records are also available, with less frequent updates, as a DarwinCore occurrence dataset via the Global Biodiversity Information Facility (<https://www.gbif.org/dataset/a267b6a7-91f9-457c-889a-481e7aa920b6>).

References

- Agassiz A (1890) Notice of *Calamocrinus diomedae*, a new stalked crinoid from the Galapagos, dredged by the U.S. Fish Commission Steamer "Albatross," Lieut.-Commander Z.L. Tanner, U.S.N., commanding. Bulletin of the Museum of Comparative Zoology at Harvard College 20: 165–167. <https://www.biodiversitylibrary.org/item/26399> [accessed on 20 May 2020]
- Agassiz A, Clark HL (1909) Hawaiian and other Pacific Echini. The Echinothuridae. Memoirs of the Museum of Comparative Zoology at Harvard College 34: 139–203. <https://doi.org/10.5962/bhl.title.42203>
- Aguado MT, Rouse GW (2011) Nautiliniellidae (Annelida) from Costa Rican cold seeps and a western Pacific hydrothermal vent, with description of four new species. Systematics and Biodiversity 9: 109–131. <https://doi.org/10.1080/14772000.2011.569033>

- Aguado MT, San Martín G, Siddall ME (2012) Systematics and evolution of syllids (Annelida, Syllidae). *Cladistics* 28: 234–250. <https://doi.org/10.1111/j.1096-0031.2011.00377.x>
- Ahyong ST, Andreakis N, Taylor J (2011) Mitochondrial phylogeny of the deep-sea squat lobsters, Munidopsidae (Galatheaidea). *Zoologischer Anzeiger* 250: 367–377. <https://doi.org/10.1016/j.jcz.2011.06.005>
- Alvarado JJ, Solís-Marín FA (Eds) (2013) Echinoderm Research and Diversity in Latin America. Springer Berlin, Heidelberg. <https://doi.org/10.1007/978-3-642-20051-9>
- Alvarado JJ, Solís-Marín FA, Ahearn CG (2010) Echinoderm (Echinodermata) diversity in the Pacific coast of Central America. *Marine Biodiversity* 40: 45–56. <https://doi.org/10.1007/s12526-009-0032-5>
- Alvarado JJ, Cortés J, Esquivel MF, Salas E (2012) Costa Rica's marine protected areas: Status and perspectives. *Revista de Biología Tropical* 60: 129–142. <https://doi.org/10.15517/rbt.v60i1.2657>
- Alvarado JJ, Chacón-Monge JL, Solís-Marín FA, Pineda-Enríquez T, Caballero-Ochoa AA, Rivera SS, Chaves RR (2017) Echinoderms from the Museum of Zoology from the Universidad de Costa Rica. *Revista de Biología Tropical* 65 (Supplement 1): S272–S287. <https://doi.org/10.15517/rbt.v65i1-1.31695>
- Alvarado JJ, Chacón-Monge JL, Azofeifa-Solano JC, Cortés J (2022) Diversity of deep-sea echinoderms from Costa Rica. *Frontiers in Marine Science* 9: 918878. <https://doi.org/10.3389/fmars.2022.918878>
- Amon DJ, Gobin J, Van Dover CL, Levin LA, Marsh L, Raineault NA (2017) Characterization of methane-seep communities in a deep-sea area designated for oil and natural gas exploitation off Trinidad and Tobago. *Frontiers in Marine Science* 4: 342. <https://doi.org/10.3389/fmars.2017.00342>
- Araya JF, Aliaga JA, Araya ME (2018) First record of *Lillipathes ritamariae* Opresko and Breedy, 2010 (Cnidaria: Antipatharia) from the southeastern Pacific Ocean. *Marine Biodiversity* 48: 1601–1605. <https://doi.org/10.1007/s12526-016-0591-1>
- Arguelles J, Larrivière P, Thatje S, Pérez M (2020) King crabs of Peruvian waters during 2003–2004: New insights. In: Hendrickx ME (Ed.) *Deep-Sea Pycnogonids and Crustaceans of the Americas*. Springer, Cham, 375–394. https://doi.org/10.1007/978-3-030-58410-8_16
- Arndt A, Marquez C, Lambert P, Smith MJ (1996) Molecular phylogeny of eastern Pacific sea cucumbers (Echinodermata: Holothuroidea) based on mitochondrial DNA sequence. *Molecular Phylogenetics and Evolution* 6: 425–437. <https://doi.org/10.1006/mpev.1996.0091>
- Arnés-Urgellés C, Buglass S, Ahyong ST, Salinas-de-León P, Wicksten MK, Marsh L (2020) Arthropoda; Crustacea; Decapoda of deep-sea volcanic habitats of the Galapagos Marine Reserve, Tropical Eastern Pacific. *Biodiversity Data Journal* 8: 1–48. <https://doi.org/10.3897/BDJ.8.e54482>
- Ashford OS, Guan S, Capone D, Rigney K, Rowley K, Cordes EE, Cortés J, Rouse GW, Mendoza GF, Sweetman AK, Levin LA (2021a) Relationships between biodiversity and ecosystem functioning proxies strengthen when approaching chemosynthetic deep-sea methane seeps. *Proceedings of the Royal Society B: Biological Sciences* 288: 20210950. <https://doi.org/10.1098/rspb.2021.0950>
- Ashford OS, Guan S, Capone D, Rigney K, Rowley K, Orphan V, Mullin SW, Dawson KS, Cortés J, Rouse GW, Mendoza GF, Lee RW, Cordes EE, Levin LA (2021b) A chemosynthetic ecotone—“chemotone”—in the sediments surrounding deep-sea methane seeps. *Limnology and Oceanography* 66: 1687–1702. <https://doi.org/10.1002/lno.11713>

- Audzijonyte A, Krylova EM, Sahling H, Vrijenhoek RC (2012) Molecular taxonomy reveals broad trans-oceanic distributions and high species diversity of deep-sea clams (Bivalvia: Vesicomyidae: Pliocardiinae) in chemosynthetic environments. *Systematics and Biodiversity* 10: 403–415. <https://doi.org/10.1080/14772000.2012.744112>
- Azofeifa-Solano JC, Cortés J (2020) Diversity and biology of deep-water crustaceans in Costa Rica. In: Hendrickx ME (Ed.) *Deep-Sea Pycnogonids and Crustaceans of the Americas*. Springer, Cham, 633–658. https://doi.org/10.1007/978-3-030-58410-8_24
- Baba K (2005) Deep-sea chirostylid and galatheid crustaceans (Decapoda: Anomura) from the Indo-Pacific, with a list of species. In: Wolff T (Ed.) *Galathea Report*. Apollo Books, Stenstrup, Denmark, 1–317.
- Baba K, Haig J (1990) A new species of chirostylid crustacean (Decapoda, Anomura) from off the west coast of North America. *Proceedings of The Biological Society of Washington* 103: 854–860. <https://www.biodiversitylibrary.org/page/34592319> [accessed on 15 April 2021]
- Baba K, Wicksten MK (2019) Chirostyloidean squat lobsters (Crustacea: Decapoda: Anomura) from the Galapagos Islands. *Zootaxa* 4564: 391–421. <https://doi.org/10.11646/zootaxa.4564.2.5>
- Baba K, Macpherson E, Poore GCB, Ah Yong S, Bermudez A, Cabezas P, Lin C-W, Nizinski M, Rodrigues C, Schnabel KE (2008) Catalogue of squat lobsters of the world (Crustacea: Decapoda: Anomura—families Chirostylidae, Galatheididae and Kiwaidae). *Zootaxa* 1905: 1–220. <https://doi.org/10.11646/zootaxa.1905.1.1>
- Bachraty C, Legendre P, Desbruyères D (2009) Biogeographic relationships among deep-sea hydrothermal vent faunas at global scale. *Deep Sea Research Part I: Oceanographic Research Papers* 56: 1371–1378. <https://doi.org/10.1016/j.dsr.2009.01.009>
- Baco AR, Smith CR (2003) High species richness in deep-sea chemoautotrophic whale skeleton communities. *Marine Ecology Progress Series* 260: 109–114. <https://doi.org/10.3354/meps260109>
- Báez P, Soto R (1997) *Paracrangon areolata* Faxon, 1893 un camarón nuevo para Chile (Decapoda, Crangonidae). *Investigaciones Marinas* 25: 277–280. <https://doi.org/10.4067/S0717-71781997002500020>
- Bailey JV, Salman V, Rouse GW, Schulz-Vogt HN, Levin LA, Orphan VJ (2011) Dimorphism in methane seep-dwelling ecotypes of the largest known bacteria. *The ISME Journal* 5: 1926–1935. <https://doi.org/10.1038/ismej.2011.66>
- Baird W (1869) Remarks on several genera of *Annelides*, belonging to the group Eunicea, with a notice of such species as are contained in the collection of the British Museum, and a description of some others hitherto undescribed. *Journal of the Linnean Society of London. Zoology* 10: 341–361. <https://doi.org/10.1111/j.1096-3642.1869.tb00665.x>
- Baker AN (2016) An illustrated catalogue of type specimens of the bathyal brittlestar genera *Ophiomusium* Lyman and *Ophiosphalma* H. L. Clark (Echinodermata: Ophiuroidea). *Zootaxa* 4097: 1–40. <https://doi.org/10.11646/zootaxa.4097.1.1>
- Ballesteros JA, Setton EVW, Santibáñez-López CE, Arango CP, Brenneis G, Brix S, Corbett KF, Cano-Sánchez E, Dandouch M, Dilly GF, Eleaume MP, Gainett G, Gallut C, McAtee S, McIntyre L, Moran AL, Moran R, López-González PJ, Scholtz G, Williamson C, Woods HA, Zehms JT, Wheeler WC, Sharma PP (2021) Phylogenomic resolution of sea spider diversification through integration of multiple data classes. *Molecular Biology and Evolution* 38: 686–701. <https://doi.org/10.1093/molbev/msaa228>
- Banse K (1980) Terebellidae (Polychaeta) from the Northeast Pacific Ocean. *Canadian Journal of Fisheries and Aquatic Sciences* 37: 20–40. <https://doi.org/10.1139/f80-003>

- Barnard JL (1967) Bathyal and abyssal Gammaridean Amphipoda of Cedros Trench, Baja California. *Bulletin of the United States National Museum* 260: 1–205. <https://doi.org/10.5479/si.03629236.260.1>
- Barry JP, Kochevar RE (1999) *Calypptogena diagonalis*, a new vesicomysid bivalve from subduction zone cold seeps in the eastern North Pacific. *The Veliger* 42: 117–123. <https://www.biodiversitylibrary.org/item/134364%0AArticle/Chapter> [accessed on 13 March 2021]
- Beaulieu SE, Baker ET, German CR, Maffei A (2013) An authoritative global database for active submarine hydrothermal vent fields. *Geochemistry, Geophysics, Geosystems* 14: 4892–4905. <https://doi.org/10.1002/2013GC004998>
- Bergmeier FS, Haszprunar G, Brandt A, Saito H, Kano Y, Jörger KM (2019) Of basins, plains, and trenches: Systematics and distribution of Solenogastres (Mollusca, Aplacophora) in the Northwest Pacific. *Progress in Oceanography* 178: 102187. <https://doi.org/10.1016/j.pocean.2019.102187>
- Bergquist DC, Urcuyo IA, Fisher CR (2002) Establishment and persistence of seep vestimentiferan aggregations on the upper Louisiana slope of the Gulf of Mexico. *Marine Ecology Progress Series* 241: 89–98. <https://doi.org/10.3354/meps241089>
- Bergquist DC, Ward T, Cordes EE, McNelis T, Howlett S, Kosoff R, Hourdez S, Carney R, Fisher CR (2003) Community structure of vestimentiferan-generated habitat islands from Gulf of Mexico cold seeps. *Journal of Experimental Marine Biology and Ecology* 289: 197–222. [https://doi.org/10.1016/S0022-0981\(03\)00046-7](https://doi.org/10.1016/S0022-0981(03)00046-7)
- Berman G, Johnson S, Seid C, Vrijenhoek R, Rouse G (2023) Range extensions of Pacific bone-eating worms (Annelida, Siboglinidae, *Osedax*). *Biodiversity Data Journal* 11: e102803. <https://doi.org/10.3897/BDJ.11.e102803>
- Bernardino AF, Levin LA, Thurber AR, Smith CR (2012) Comparative composition, diversity and trophic ecology of sediment macrofauna at vents, seeps and organic falls. *PLoS ONE* 7(4): e33515. <https://doi.org/10.1371/journal.pone.0033515>
- Bettters MJ, Cordes EE (2024) New records of *Provanna* (Gastropoda, Provannidae) from the Costa Rica Margin and an identification key for the genus. *ZooKeys* 1189: 1–32. <https://doi.org/10.3897/zookeys.1189.109734>
- Bettters MJ, Cortés J, Cordes EE (2024) New species and records of limpets (Mollusca, Gastropoda) from the Pacific Costa Rica Margin. *ZooKeys* 1214: 281–324. <https://doi.org/10.3897/zookeys.1214.128594>
- Bettters M, Stabbins A, Keller A, Cordes E (2023) Biogeography and depth partitioning in deep-sea gastropods at the Pacific Costa Rica Margin. *Journal of Biogeography*: 1–13. <https://doi.org/10.1111/jbi.14722>
- Bigelow RP (1894) Report upon the Crustacea of the order Stomatopoda collected by the Steamer Albatross between 1885 and 1891, and on other specimens in the U.S. National Museum. *Proceedings of the United States National Museum* 17: 489–544. <https://doi.org/10.1126/science.9.231.776>
- Black MB, Halanych KM, Maas PAY, Hoeh WR, Hashimoto J, Desbruyères D, Lutz RA, Vrijenhoek RC (1997) Molecular systematics of vestimentiferan tubeworms from hydrothermal vents and cold-water seeps. *Marine Biology* 130: 141–149. <https://doi.org/10.1007/s002270050233>
- Black MB, Trivedi A, Maas PAY, Lutz RA, Vrijenhoek RC (1998) Population genetics and biogeography of vestimentiferan tube worms. *Deep Sea Research Part II: Topical Studies in Oceanography* 45: 365–382. [https://doi.org/10.1016/S0967-0645\(97\)00076-3](https://doi.org/10.1016/S0967-0645(97)00076-3)
- Blake JA (1985) Polychaeta from the vicinity of deep-sea geothermal vents in the eastern Pacific. I: Euphrosinidae, Phyllodocidae, Hesionidae, Nereididae, Glyceridae,

- Dorvilleidae, Orbiniidae, and Maldanidae. *Bulletin of the Biological Society of Washington* 6: 67–101.
- Blake JA (1996) Family Paraonidae. In: Blake JA, Hilbig B, Scott PH (Eds) *Taxonomic Atlas of the Benthic Fauna of the Santa Maria Basin and Western Santa Barbara Channel. Volume 6. The Annelida Part 3 - Polychaeta: Orbiniidae to Cossuridae*. Santa Barbara Museum of Natural History, CA, USA, 27–70.
- Blake JA, Hilbig B (1990) Polychaeta from the vicinity of deep-sea hydrothermal vents in the eastern Pacific. II. New species and records from the Juan de Fuca and Explorer Ridge systems. *Pacific Science* 44: 219–253. <https://scholarspace.manoa.hawaii.edu/server/api/core/bitstreams/deff09af-c55c-48db-97cd-c1d5edce7f0d/content> [accessed on 4 May 2020]
- Blake JA, Maciolek NJ (1992) Polychaeta from deep-sea hydrothermal vents in the eastern Pacific. III. A new genus and two new species of Spionidae from the Guaymas Basin and Juan de Fuca Ridge with comments on a related species from the western North Atlantic. *Proceedings of the Biological Society of Washington* 105: 723–732. <https://biodiversitylibrary.org/page/35607908> [accessed on 2 April 2021]
- Blum S, Fong J (2016a) *Araeosoma leptaleum* A. Agassiz & H.L. Clark, 1909 occurrence 609472526. CAS Invertebrate Zoology (IZ). Version 14.2. California Academy of Sciences occurrence dataset. <https://doi.org/10.15468/tiac99>
- Blum S, Fong J (2016b) *Araeosoma leptaleum* A. Agassiz & H.L. Clark, 1909 occurrence 609475715. CAS Invertebrate Zoology (IZ). Version 14.2. California Academy of Sciences occurrence dataset. <https://doi.org/10.15468/tiac99>
- Blum S, Fong J (2016c) *Delectopecten vancouverensis* (Whiteaves, 1893) occurrence 609414722. CAS Invertebrate Zoology (IZ). Version 14.2. California Academy of Sciences occurrence dataset. <https://doi.org/10.15468/tiac99>
- Boetius A, Ravensschlag K, Schubert CJ, Rickert D, Widdel F, Gieseke A, Amann R, Jørgensen BB, Witte U, Pfannkuche O (2000) A marine microbial consortium apparently mediating anaerobic oxidation of methane. *Nature* 407: 623–626. <https://doi.org/10.1038/35036572>
- Bohrmann G, Heeschen K, Jung C, Weinrebe W, Baranov B, Atrice Cailleau B, Heath R, Hü V, Hort M, Masson D, Trummer I (2002) Widespread fluid expulsion along the seafloor of the Costa Rica convergent margin. *Terra Nova* 14: 69–79. <https://doi.org/10.1046/j.1365-3121.2002.00400.x>
- Boratyn GM, Camacho C, Cooper PS, Coulouris G, Fong A, Ma N, Madden TL, Matten WT, McGinnis SD, Merezhuk Y, Raytselis Y, Sayers EW, Tao T, Ye J, Zaretskaya I (2013) BLAST: a more efficient report with usability improvements. *Nucleic Acids Research* 41: W29–W33. <https://doi.org/10.1093/nar/gkt282>
- Borda E, Kudenov JD, Chevaldonné P, Blake JA, Desbruyères D, Fabri MC, Hourdez S, Pleijel F, Shank TM, Wilson NG, Schulze A, Rouse GW (2013) Cryptic species of *Archinome* (Annelida: Amphinomida) from vents and seeps. *Proceedings of the Royal Society B: Biological Sciences* 280. <https://doi.org/10.1098/rspb.2013.1876>
- Bouchet P, Warén A (1986) Revision of the northeast Atlantic bathyal and abyssal Aclididae, Eulimidae, Epitoniidae (Mollusca, Gastropoda). *Bollettino Malacologico Suppl.* 2: 297–576. <https://doi.org/10.5962/bhl.title.140762>
- Bouchet P, Rocroi JP, Hausdorf B, Kaim A, Kano Y, Nützel A, Parkhaev P, Schrödl M, Strong EE (2017) Revised classification, nomenclator and typification of gastropod and monoplacophoran families. *Malacologia* 61: 1–526. <https://doi.org/10.4002/040.061.0201>

- Bradbury MG (2003) Family Ogcocephalidae Jordan 1895 batfishes. California Academy of Sciences Annotated Checklists of Fishes 17: 1–17. <https://www.calacademy.org/scientists/annotated-checklists-of-fishes> [accessed on 6 May 2021]
- Breedy O, van Ofwegen LP, Vargas S (2012) A new family of soft corals (Anthozoa, Octocorallia, Alcyonacea) from the aphotic tropical eastern Pacific waters revealed by integrative taxonomy. *Systematics and Biodiversity* 10: 351–359. <https://doi.org/10.1080/14772000.2012.707694>
- Breedy O, Rouse GW, Stabbins A, Cortés J, Cordes EE (2019) New records of *Swiftia* (Cnidaria, Anthozoa, Octocorallia) from off the Pacific Costa Rican margin, including a new species from methane seeps. *Zootaxa* 4671: 407–419. <https://doi.org/10.11646/zootaxa.4671.3.6>
- Bribiesca-Contreras G, Dahlgren TG, Amon DJ, Cairns S, Drennan R, Durden JM, Eléaume MP, Hosie AM, Kremenetskaia A, McQuaid K, O'Hara TD, Rabone M, Simon-Lledó E, Smith CR, Watling L, Wiklund H, Glover AG (2022) Benthic megafauna of the western Clarion-Clipperton Zone, Pacific Ocean. *ZooKeys* 1113: 1–110. <https://doi.org/10.3897/zookeys.1113.82172>
- Bright M, Lallier F (2010) The biology of vestimentiferan tubeworms. In: Gibson RN, Atkinson RJA, Gordon JDM (Eds), *Oceanography and Marine Biology: An Annual Review*. Chapman and Hall/CRC, 213–265. <https://doi.org/10.1201/EBK1439821169>
- Brzechffa C, Goffredi SK (2021) Contrasting influences on bacterial symbiont specificity by co-occurring deep-sea mussels and tubeworms. *Environmental Microbiology Reports* 13: 104–111. <https://doi.org/10.1111/1758-2229.12909>
- Buckner JC, Sanders RC, Faircloth BC, Chakrabarty P (2021) The critical importance of vouchers in genomics. *eLife* 10: e68264. <https://doi.org/10.7554/eLife.68264>
- Cairns SD (1991) A revision of the ahermatypic Scleractinia of the Galápagos and Cocos Islands. *Smithsonian Contributions to Zoology*: 1–32. <https://doi.org/10.5479/si.00810282.504>
- Cairns SD (2006) New records of azooxanthellate Scleractinia from the Hawaiian Islands. *Bishop Museum Occasional Papers* 87: 45–53.
- Camp DK, Kuck HG (1990) Additional records of stomatopod crustaceans from Isla del Coco and Golfo de Papagayo, East Pacific Ocean. *Proceedings of the Biological Society of Washington* 103: 847–853. <https://www.biodiversitylibrary.org/item/107575> [accessed on 12 June 2020]
- Cannon JT, Swalla BJ, Halanych KM (2013) Hemichordate molecular phylogeny reveals a novel cold-water clade of harrimaniid acorn worms. *The Biological Bulletin* 225: 194–204. <https://doi.org/10.1086/BBLv225n3p194>
- Capa M, Aguado MT, Bakken T (2016) Phylogenetic hypothesis of Sphaerodoridae Malmgren, 1867 (Annelida) and its position within Phyllodocida. *Cladistics* 32: 335–350. <https://doi.org/10.1111/cla.12134>
- Capa M, Nygren A, Parapar J, Bakken T, Meißner K, Moreira J (2019) Systematic re-structure and new species of Sphaerodoridae (Annelida) after morphological revision and molecular phylogenetic analyses of the North East Atlantic fauna. *ZooKeys* 845: 1–97. <https://doi.org/10.3897/zookeys.845.32428>
- Cardoso P, Erwin TL, Borges PAV, New TR (2011) The seven impediments in invertebrate conservation and how to overcome them. *Biological Conservation* 144: 2647–2655. <https://doi.org/10.1016/j.biocon.2011.07.024>
- Carney RS (1994) Consideration of the oasis analogy for chemosynthetic communities at Gulf of Mexico hydrocarbon vents. *Geo-Marine Letters* 14: 149–159. <https://doi.org/10.1007/BF01203726>

- Carr CM, Hardy SM, Brown TM, Macdonald TA, Hebert PDN (2011) A tri-oceanic perspective: DNA barcoding reveals geographic structure and cryptic diversity in Canadian polychaetes. *PLoS ONE* 6: e22232. <https://doi.org/10.1371/journal.pone.0022232>
- Case DH, Pasulka AL, Marlow JJ, Grupe BM, Levin LA, Orphan VJ (2015) Methane seep carbonates host distinct, diverse, and dynamic microbial assemblages. *mBio* 6: e01348-15. <https://doi.org/10.1128/mBio.01348-15>
- Chan BKK, Dreyer N, Gale AS, Glenner H, Ewers-Saucedo C, Pérez-Losada M, Kolbasov GA, Crandall KA, Høeg JT (2021) The evolutionary diversity of barnacles, with an updated classification of fossil and living forms. *Zoological Journal of the Linnean Society* 193: 789–846. <https://doi.org/10.1093/zoolinnean/zlaa160>
- Chapman ASA, Beaulieu SE, Colaço A, Gebruk AV, Hilario A, Kihara TC, Ramirez-Llodra E, Sarrazin J, Tunnicliffe V, Amon DJ, Baker MC, Boschen-Rose RE, Chen C, Cooper IJ, Copley JT, Corbari L, Cordes EE, Cuvelier D, Duperron S, Du Preez C, Gollner S, Horton T, Hourdez S, Krylova EM, Linse K, LokaBharathi PA, Marsh L, Matabos M, Mills SW, Mullineaux LS, Rapp HT, Reid WDK, Rybakova E, Tresa TR, Southgate SJ, Stöhr S, Turner PJ, Watanabe HK, Yasuhara M, Bates AE (2019) sFDvent: A global trait database for deep-sea hydrothermal-vent fauna. *Global Ecology and Biogeography* 28: 1538–1551. <https://doi.org/10.1111/geb.12975>
- Chen C, Watanabe HK, Sasaki T (2019a) Four new deep-sea provannid snails (Gastropoda: Abysochrysoidea) discovered from hydrocarbon seep and hydrothermal vents in Japan. *Royal Society Open Science* 6(7): 190393. <https://doi.org/10.1098/rsos.190393>
- Chen C, Watanabe HK, Nagai Y, Toyofuku T, Xu T, Sun J, Qiu JW, Sasaki T (2019b) Complex factors shape phenotypic variation in deep-sea limpets. *Biology Letters* 15: 1–7. <https://doi.org/10.1098/rsbl.2019.0504>
- Chen C, Zhong Z, Qiu JW, Sun J (2023) A new *Paralepetopsis* limpet from a South China Sea seep hints at a paraphyletic Neolepetopsidae. *Zoological Studies* 62: e26. <https://doi.org/10.6620/ZS.2023.62-26>
- Chen C, Hookabe N, Komatsu H (2024a) Faunal community of a new hot vent field on the Amami Rift. *Journal of the Marine Biological Association of the United Kingdom* 104: e39. <https://doi.org/10.1017/S0025315424000304>
- Chen C, Jamieson JW, Tunnicliffe V (2024b) Hydrothermal vent fauna of the Galápagos Rift: updated species list with new records. *Marine Biodiversity* 54: 16. <https://doi.org/10.1007/s12526-024-01408-w>
- Chen C, Watanabe HK, Sawada H, Iwamoto H, Takai K (2024c) Serpentinite-hosted chemosynthetic community of South Chamorro Seamount, Mariana Forearc. *Marine Ecology* 45(4): e12808. <https://doi.org/10.1111/maec.12808>
- Christodoulou M, O'Hara TD, Hugall AF, Arbizu PM (2019) Dark ophiuroid biodiversity in a prospective abyssal mine field. *Current Biology* 29: 3909–3912.e3. <https://doi.org/10.1016/j.cub.2019.09.012>
- Christodoulou M, O'Hara T, Hugall AF, Khodami S, Rodrigues CF, Hilario A, Vink A, Arbizu PM (2020) Unexpected high abyssal ophiuroid diversity in polymetallic nodule fields of the northeast Pacific Ocean and implications for conservation. *Biogeosciences* 17: 1845–1876. <https://doi.org/10.5194/bg-17-1845-2020>
- Clark AH (1907) Descriptions of new species of recent unstalked crinoids from the North Pacific Ocean. *Proceedings of the United States National Museum* 33: 69–84. <https://doi.org/10.5479/si.00963801.33-1559.69>
- Clark HL (1911) North Pacific ophiurans in the collection of the United States National Museum. *Bulletin of the United States National Museum* 75: 1–302. <https://doi.org/10.5479/si.03629236.75.1>

- Clark HL (1912) Hawaiian and other Pacific Echini. The Pedinidae, Phymosomatidae, Stomopneustidae, and Echinometridae. *Memoirs of the Museum of Comparative Zoology at Harvard College* 34: 205–383. <https://doi.org/10.5962/bhl.title.42203>
- Clark HL (1913) Echinoderms from Lower California, with descriptions of new species. *Bulletin of the American Museum of Natural History* 32: 185–236. <https://doi.org/10.5962/bhl.title.1734>
- Clark AH (1915) A monograph of the existing crinoids, volume 1: The comatulids, part 2. *Bulletin of the United States National Museum* 82: 1–406. <https://doi.org/10.5962/bhl.title.1326>
- Clark AH (1916a) Six new starfishes from the Gulf of California and adjacent waters. *Proceedings of the Biological Society of Washington* 29: 51–62. <https://www.biodiversitylibrary.org/page/3336447> [accessed on 13 May 2021]
- Clark HL (1916b) Report on the sea-lilies, starfishes, brittle-stars and sea-urchins obtained by the F.I.S. “Endeavour” on the coasts of Queensland, New South Wales, Tasmania, Victoria, South Australia, and Western Australia. In: Dannevig HC (Ed.) *Biological results of the fishing experiments carried on by the “Endeavour”, v. 4, pt. 1*. Commonwealth of Australia Department of Trade and Customs, Sydney, 1–123. <https://doi.org/10.5962/bhl.title.13854>
- Clark HL (1917) Reports on the scientific results of the expedition to the eastern tropical Pacific in charge of Alexander Agassiz by the U.S. Fish Commission steamer Albatross, from October, 1904, to March, 1905, Lieut. Commander L.M. Garrett, U.S.N., commanding. XXX. Ophiuroidea. *Bulletin of the Museum of Comparative Zoology at Harvard College* 61: 429–453. <https://www.biodiversitylibrary.org/part/11461> [accessed on 25 April 2022]
- Clark K, Karsch-Mizrachi I, Lipman DJ, Ostell J, Sayers EW (2016) GenBank. *Nucleic Acids Research* 44: D67–D72. <https://doi.org/10.1093/nar/gkv1276>
- Clement M, Snell Q, Walker P, Posada D, Crandall K (2002) TCS: Estimating gene genealogies. In: *Proceedings of the 6th International Parallel and Distributed Processing Symposium*. IEEE, Fort Lauderdale, FL. <https://doi.org/10.1109/IPDPS.2002.1016585>
- Coan EV, Valentich-Scott P (2012) *Bivalve Seashells of Tropical West America: Marine Bivalve Mollusks from Baja California to Northern Perú*. 1st edn. Santa Barbara Museum of Natural History, CA, USA.
- Coan EV, Valentich-Scott P, Bernard FR (2000) *Bivalve Seashells of Western North America: Marine Bivalve Mollusks from Arctic Alaska to Baja California*. 1st edn. Santa Barbara Museum of Natural History, CA, USA.
- Copilaş-Ciocianu D, Borko Š, Fišer C (2020) The late blooming amphipods: Global change promoted post-Jurassic ecological radiation despite Palaeozoic origin. *Molecular Phylogenetics and Evolution* 143: 106664. <https://doi.org/10.1016/j.ympev.2019.106664>
- Cordes EE, Becker EL, Hourdez S, Fisher CR (2010) Influence of foundation species, depth, and location on diversity and community composition at Gulf of Mexico lower-slope cold seeps. *Deep Sea Research Part II: Topical Studies in Oceanography* 57: 1870–1881. <https://doi.org/10.1016/j.dsr2.2010.05.010>
- Cortés J (2012) Marine biodiversity of an Eastern Tropical Pacific oceanic island, Isla del Coco, Costa Rica. *Revista de Biología Tropical* 60 (Supplement 3): 131–185. <https://doi.org/10.15517/rbt.v60i3.28356>
- Cortés J, Wehrtmann IS (2009) Diversity of marine habitats of the Caribbean and Pacific of Costa Rica. In: Wehrtmann IS, Cortés J (Eds), *Marine Biodiversity of Costa Rica, Central America*. Monographiae Biologicae. Springer, Dordrecht, 1–45. https://doi.org/10.1007/978-1-4020-8278-8_1

- Costa FO, DeWaard JR, Boutillier J, Ratnasingham S, Dooh RT, Hajibabaei M, Hebert PDN (2007) Biological identifications through DNA barcodes: The case of the Crustacea. *Canadian Journal of Fisheries and Aquatic Sciences* 64: 272–295. <https://doi.org/10.1139/f07-008>
- Cowart DA, Huang C, Arnaud-Haond S, Carney SL, Fisher CR, Schaeffer SW (2013) Restriction to large-scale gene flow vs. regional panmixia among cold seep *Escarpia* spp. (Polychaeta, Siboglinidae). *Molecular Ecology* 22: 4147–4162. <https://doi.org/10.1111/mec.12379>
- Dahlgren TG, Wiklund H, Rabone M, Amon DJ, Ikebe C, Watling L, Smith CR, Glover AG (2016) Abyssal fauna of the UK-1 polymetallic nodule exploration area, Clarion-Cliperton Zone, central Pacific Ocean: Cnidaria. *Biodiversity Data Journal* 4: 1–31. <https://doi.org/10.3897/BDJ.4.e9277>
- Dall WH (1891) Scientific results of explorations by the U. S. Fish Commission steamer Albatross. No. XX.—On some new or interesting west American shells obtained from the dredgings of the U. S. Fish Commission steamer Albatross in 1888, and from other sources. *Proceedings of the United States National Museum* 14: 173–191. <https://doi.org/10.5479/si.00963801.14-849.173>
- Dall WH (1896) Diagnoses of new species of molluscs from the west coast of America. *Proceedings of the United States National Museum* 18: 7–20. <https://doi.org/10.5479/si.00963801.1034>
- Dall WH (1902) Illustrations and descriptions of new, unfigured, or imperfectly known shells, chiefly American, in the U.S. National Museum. *Proceedings of the United States National Museum* 24: 499–566. <https://doi.org/10.5479/si.00963801.24-1264.499>
- Dall WH (1908) Reports on the dredging operations off the west coast of Central America to the Galapagos, to the west coast of Mexico, and in the Gulf of California, in charge of Alexander Agassiz, carried on by the U.S. Fish Commission steamer “Albatross,” during 1891, Lieut.-Commander Z.L. Tanner, U.S.N., commanding. XXXVII. Reports on the scientific results of the expedition to the eastern tropical Pacific, in charge of Alexander Agassiz, by the U.S. Fish Commission steamer “Albatross”, from October, 1904 to March, 1905, Lieut.-Commander L.M. Garrett, U.S.N., commanding. XIV. The Mollusca and Brachiopoda. *Bulletin of the Museum of Comparative Zoology at Harvard College* 43: 205–487. <https://doi.org/10.5962/bhl.title.46539>
- Darwin Core Maintenance Group (2021) Darwin Core Quick Reference Guide. Biodiversity Information Standards (TDWG). <https://dwc.tdwg.org/terms/> [accessed on 9 June 2024]
- Deagle BE, Bax N, Hewitt CL, Patil JG (2003) Development and evaluation of a PCR-based test for detection of *Asterias* (Echinodermata: Asteroidea) larvae in Australian plankton samples from ballast water. *Marine and Freshwater Research* 54: 709–719. <https://doi.org/10.1071/MF03031>
- Decker C, Olu K, Cunha RL, Arnaud-Haond S (2012) Phylogeny and diversification patterns among vesicomyid bivalves. *PLoS ONE* 7: e33359. <https://doi.org/10.1371/journal.pone.0033359>
- Dekas AE, Chadwick GL, Bowles MW, Joye SB, Orphan VJ (2014) Spatial distribution of nitrogen fixation in methane seep sediment and the role of the ANME archaea. *Environmental Microbiology* 16: 3012–3029. <https://doi.org/10.1111/1462-2920.12247>
- Desbruyères D, Segonzac M, Bright M (Eds) (2006) 18 Handbook of Deep-Sea Hydrothermal Vent Fauna. 2nd edn. Biologiezentrum der Oberösterreichischen Landesmuseen, Linz, Austria. https://www.zobodat.at/publikation_volumes.php?id=23702 [accessed on 4 March 2021]

- Downey ME (1968) Catalog of recent echinoid type specimens in the U.S. National Museum, Smithsonian Institution, and the Museum of Comparative Zoology, Harvard University. Bulletin of the United States National Museum 264: 1–99. <https://doi.org/10.5479/si.03629236.264>
- Drennan R, Wiklund H, Rouse GW, Georgieva MN, Wu X, Kobayashi G, Yoshino K, Glover AG (2019) Taxonomy and phylogeny of mud owls (Annelida: Sternaspidae), including a new synonymy and new records from the Southern Ocean, North East Atlantic Ocean and Pacific Ocean: challenges in morphological delimitation. Marine Biodiversity 49: 2659–2697. <https://doi.org/10.1007/s12526-019-00998-0>
- Duchassaing P, Michelotti J (1864) Supplément au mémoire sur les coralliaires des Antilles. Memorie della Reale Accademia della Scienze di Torino 2: 1–112. <https://doi.org/10.5962/bhl.title.105196>
- Dunn CW, Giribet G, Edgecombe GD, Hejnol A (2014) Animal phylogeny and its evolutionary implications. Annual Review of Ecology, Evolution, and Systematics 45: 371–395. <https://doi.org/10.1146/annurev-ecolsys-120213-091627>
- Edgecombe GD, Giribet G, Wheeler WC (2002) Phylogeny of Henicopidae (Chilopoda: Lithobiomorpha): A combined analysis of morphology and five molecular loci. Systematic Entomology 27: 31–64. <https://doi.org/10.1046/j.0307-6970.2001.00163.x>
- Eilertsen MH, Georgieva MN, Kongsrud JA, Linse K, Wiklund H, Glover AG, Rapp HT (2018) Genetic connectivity from the Arctic to the Antarctic: *Sclerolinum contortum* and *Nicomache lokii* (Annelida) are both widespread in reducing environments. Scientific Reports 8: 4810. <https://doi.org/10.1038/s41598-018-23076-0>
- Eilertsen MH, Kongsrud JA, Tandberg AHS, Alvestad T, Budaeva N, Martell L, Ramalho SP, Falkenhaug T, Huys R, Oug E, Bakken T, Høisæter T, Rauch C, Carvalho FC, Savchenko AS, Ulvatn T, Kongshavn K, Berntsen CM, Olsen BR, Pedersen RB (2024) Diversity, habitat endemism and trophic ecology of the fauna of Loki's Castle vent field on the Arctic Mid-Ocean Ridge. Scientific Reports 14: 103. <https://doi.org/10.1038/s41598-023-46434-z>
- Faxon W (1893) Reports on the dredging operations off the west coast of Central America to the Galapagos, to the west coast of Mexico, and in the Gulf of California, in charge of Alexander Agassiz, carried on by the U.S. Fish Commission steamer "Albatross", during 1891, Lieut. Commander Z.L. Tanner, U.S.N., commanding. VI. Preliminary descriptions of new species of Crustacea. Bulletin of the Museum of Comparative Zoology at Harvard College 24: 149–220. <https://www.marinespecies.org/aphia.php?p=sourceget&id=130134> [accessed on 6 May 2020]
- Faxon W (1895) Reports on an exploration off the west coasts of Mexico, Central and South America, and off the Galapagos Islands, in charge of Alexander Agassiz, by the U.S. Fish Commission Steamer "Albatross," during 1891, Lieut.-Commander Z. L. Tanner, U.S.N. commanding. XV. The stalk-eyed Crustacea. Memoirs of the Museum of Comparative Zoology at Harvard College 18: 1–292. <https://doi.org/10.5962/bhl.title.14033>
- Feldman RA, Shank TM, Black MB, Baco AR, Smith CR, Vrijenhoek RC (1998) Vestimentiferan on a whale fall. The Biological Bulletin 194: 116–119. <https://doi.org/10.2307/1543041>
- Fiedler PC, Lavín MF (2006) Introduction: A review of eastern tropical Pacific oceanography. Progress in Oceanography 69: 94–100. <https://doi.org/10.1016/j.pocean.2006.03.006>
- Fischer W, Krupp F, Schneider W, Sommer C, Carpenter KE, Niem VH (Eds) (1995) Guía FAO para la Identificación de Especies para los Fines de la Pesca. Pacífico centro-oriental. Volumen I. Plantas e Invertebrados. FAO, Rome. <https://www.fao.org/3/t0851s/t0851s00.htm> [accessed on 23 April 2021]

- Fisher WK (1905) New starfishes from deep water off California and Alaska. Bulletin of the Bureau of Fisheries 24: 291–320. <http://www.biodiversitylibrary.org/page/27012292/> [accessed on 19 May 2020]
- Fisher WK (1906) New starfishes from the Pacific coast of North America. Proceedings of the Washington Academy of Sciences 8: 111–139. <https://www.jstor.org/stable/24526115> [accessed on 13 May 2020]
- Fisher WK (1911) Asteroidea of the North Pacific and adjacent waters. Part 1: Phanerozonia and Spinulosa. Bulletin of the United States National Museum 76: 1–419. <https://doi.org/10.5479/si.03629236.76.i>
- Folmer O, Black M, Hoeh W, Lutz R, Vrijenhoek R (1994) DNA primers for amplification of mitochondrial cytochrome c oxidase subunit I from diverse metazoan invertebrates. Molecular Marine Biology and Biotechnology 3: 294–299. <https://doi.org/10.1071/ZO9660275>
- Foster MW (1989) Brachiopods from the extreme South Pacific and adjacent waters. Journal of Paleontology 63: 268–301. <https://doi.org/10.1017/S0022336000019442>
- Frable BW, Seid CA, Bronson AW, Moller PR (2023) A new deep-sea eelpout of the genus *Pyrolycus* (Teleostei: Zoarcidae) associated with a hydrothermal seep on the Pacific margin of Costa Rica. Zootaxa 5230: 79–89. <https://doi.org/10.11646/zootaxa.5230.1.5>
- Fujikura K, Kojima S, Tamaki K, Maki Y, Hunt J, Okutani T (1999) The deepest chemosynthesis-based community yet discovered from the hadal zone, 7326 m deep, in the Japan Trench. Marine Ecology Progress Series 190: 17–26. <https://doi.org/10.3354/meps190017>
- Fukasawa Y, Matsumoto H, Beppu S, Fujiwara Y, Kawato M, Miyazaki J-I (2017) Molecular phylogenetic analysis of chemosymbiotic Solemyidae and Thyasiridae. Open Journal of Marine Science 7: 124–141. <https://doi.org/10.4236/ojms.2017.71010>
- Galil BS (2000) Crustacea Decapoda: Review of the genera and species of the family Polychelidae Wood-Mason, 1874. In: Crosnier A (Ed.) Résultats des Campagnes MUSORSTOM, Volume 21. Mémoires du Muséum national d'histoire naturelle, Paris, 285–387. <https://decapoda.nhm.org/pdfs/16080/16080.pdf> [accessed on 7 May 2020]
- Gallo ND, Cameron J, Hardy K, Fryer P, Bartlett DH, Levin LA (2015) Submersible- and lander-observed community patterns in the Mariana and New Britain trenches: Influence of productivity and depth on epibenthic and scavenging communities. Deep Sea Research Part I: Oceanographic Research Papers 99: 119–133. <https://doi.org/10.1016/j.dsr.2014.12.012>
- GBIF: The Global Biodiversity Information Facility (2024) What is GBIF? <https://www.gbif.org/what-is-gbif> [accessed on 3 June 2024]
- GBIF.org (2022a) *Ophiacantha moniliformis* occurrences. GBIF Occurrence Download. <https://doi.org/10.15468/dl.6v92gz>
- GBIF.org (2022b) *Ophiophrura liodisca* occurrences. GBIF Occurrence Download. <https://doi.org/10.15468/dl.3dupdc>
- GBIF.org (2024) *Ophiocreas carnosus* occurrences. GBIF Occurrence Download. <https://doi.org/10.15468/dl.vkckcr>
- Geller JB, Walton ED (2001) Breaking up and getting together: Evolution of symbiosis and cloning by fission in sea anemones (genus *Anthopleura*). Evolution 55: 1781–1794. <https://doi.org/10.1111/j.0014-3820.2001.tb00827.x>
- Geller J, Meyer C, Parker M, Hawk H (2013) Redesign of PCR primers for mitochondrial cytochrome c oxidase subunit I for marine invertebrates and application in all-taxa biotic surveys. Molecular Ecology Resources 13: 851–861. <https://doi.org/10.1111/1755-0998.12138>

- Gemeinholzer B, Rey I, Weising K, Grundmann M, Muellner AN, Zetzsche H, Droege G (2010) Chapter 7 Organizing specimen and tissue preservation in the field for subsequent molecular analyses. In: Eymann J, Degreef J, Häuser C, Monje JC, Samyn Y, Vandenspiegel D (Eds), Manual on Field Recording Techniques and Protocols for All Taxa Biodiversity Inventories. *Abc Taxa* 8: 129–157. <https://abctaxa.naturalsciences.be/pdf-8/> [accessed on 27 February 2017]
- German CR, Ramirez-Llodra E, Baker MC, Tyler PA, Baco-Taylor A, Boetius A, Bright M, de Siqueira LC, Cordes EE, Desbruyères D, Dubilier N, Fisher CR, Fujiwara Y, Gaill F, Gebruk A, Juniper K, Levin LA, Lokabharathi PA, Metaxas A, Rowden AA, Santos RS, Shank TM, Smith CR, Van Dover CL, Young CM, Warén A (2011) Deep-water chemosynthetic ecosystem research during the Census of Marine Life decade and beyond: A proposed deep-ocean road map. *PLoS ONE* 6(8): e23259. <https://doi.org/10.1371/journal.pone.0023259>
- Giribet G, Carranza S, Bagui J, Riutort M, Ribera C (1996) First molecular evidence for the existence of a Tardigrada + Arthropoda clade. *Molecular Biology and Evolution* 13: 76–84. <https://doi.org/10.1093/oxfordjournals.molbev.a025573>
- Giribet G, Carranza S, Riutort M, Baguñà J, Ribera C (1999) Internal phylogeny of the Chilopoda (Myriapoda, Arthropoda) using complete 18s rDNA and partial 28S rDNA sequences. *Philosophical Transactions of the Royal Society of London. Series B, Biological Sciences* 354: 215–222. <https://doi.org/10.1098/rstb.1999.0373>
- Glover AG, Goetze E, Dahlgren TG, Smith CR (2005) Morphology, reproductive biology and genetic structure of the whale-fall and hydrothermal vent specialist, *Bathyscuria guaymasensis* Pettibone, 1989 (Annelida: Polynoidae). *Marine Ecology* 26: 223–234. <https://doi.org/10.1111/j.1439-0485.2005.00060.x>
- Glover AG, Dahlgren TG, Wiklund H, Mohrbeck I, Smith CR (2015) An end-to-end DNA taxonomy methodology for benthic biodiversity survey in the Clarion-Clipperton Zone, central Pacific abyss. *Journal of Marine Science and Engineering* 4: 2. <https://doi.org/10.3390/jmse4010002>
- Glover AG, Wiklund H, Rabone M, Amon DJ, Smith CR, O'Hara T, Mah CL, Dahlgren TG (2016) Abyssal fauna of the UK-1 polymetallic nodule exploration claim, Clarion-Clipperton Zone, central Pacific Ocean: Echinodermata. *Biodiversity Data Journal* 4: e7251. <https://doi.org/10.3897/BDJ.4.e7251>
- Glover AG, Wiklund H, Chen C, Dahlgren TG (2018) Point of view: Managing a sustainable deep-sea 'blue economy' requires knowledge of what actually lives there. *eLife* 7: e41319. <https://doi.org/10.7554/eLife.41319>
- Goffredi SK, Hurtado LA, Hallam S, Vrijenhoek RC (2003) Evolutionary relationships of deep-sea vent and cold seep clams (Mollusca: Vesicomyidae) of the "pacific/lepta" species complex. *Marine Biology* 142: 311–320. <https://doi.org/10.1007/s00227-002-0941-3>
- Goffredi SK, Gregory A, Jones WJ, Morella NM, Sakamoto RI (2014) Ontogenetic variation in epibiont community structure in the deep-sea yeti crab, *Kiwa puravida*: convergence among crustaceans. *Molecular Ecology* 23: 1457–1472. <https://doi.org/10.1111/mec.12439>
- Goffredi SK, Johnson S, Tunnicliffe V, Caress D, Clague D, Escobar E, Lundsten L, Paduan JB, Rouse G, Salcedo DL, Soto LA, Spelz-Madero R, Zierenberg R, Vrijenhoek R (2017) Hydrothermal vent fields discovered in the southern Gulf of California clarify role of habitat in augmenting regional diversity. *Proceedings of the Royal Society B: Biological Sciences* 284: 20170817. <https://doi.org/10.1098/rspb.2017.0817>
- Goffredi SK, Tilic E, Mullin SW, Dawson KS, Keller A, Lee RW, Wu F, Levin LA, Rouse GW, Cordes EE, Orphan VJ (2020) Methanotrophic bacterial symbionts fuel dense

- populations of deep-sea feather duster worms (Sabellida, Annelida) and extend the spatial influence of methane seepage. *Science Advances* 6: eaay8562. <https://doi.org/10.1126/sciadv.aay8562>
- Goffredi SK, Motooka C, Fike DA, Gusmão LC, Tilic E, Rouse GW, Rodríguez E (2021) Mixotrophic chemosynthesis in a deep-sea anemone from hydrothermal vents in the Pescadero Basin, Gulf of California. *BMC Biology* 19: 1–18. <https://doi.org/10.1186/s12915-020-00921-1>
- Goode GB (1884) Descriptive catalogues constituting a report upon the exhibit of the fisheries and fish culture of the United States of America, made at the London Fisheries Exhibition, 1883. *Bulletin of the United States National Museum* 27: 1–1249. <https://library.si.edu/digital-library/book/bulletinunit271884unit> [accessed on 19 May 2020]
- Goto R, Monnington J, Sciberras M, Hirabayashi I, Rouse GW (2020) Phylogeny of Echiura updated, with a revised taxonomy to reflect their placement in Annelida as sister group to Capitellidae. *Invertebrate Systematics* 34: 101–111. <https://doi.org/10.1071/IS19020>
- Goy JW (1980) *Spongicoloides galapagensis*, a new shrimp representing the first record of the genus from the Pacific Ocean (Crustacea: Decapoda: Stenopodidea). *Proceedings of the Biological Society of Washington* 93: 760–770. <https://decapoda.nhm.org/pdfs/25554/25554.pdf> [accessed on 26 April 2021]
- Gracia C A, Levin LA, Zea S (2020) Meio-epifaunal wood colonization in the vicinity of methane seeps. *Marine Ecology* 41: e12573. <https://doi.org/10.1111/maec.12573>
- Granja-Fernández R, Herrero-Pérezrul MD, López-Pérez A, Hernández-Morales A, Rangel-Solís PD (2015a) A literature review of the Ophiuroidea (Echinodermata) from the Pacific coast of Mexico. *Revista de Biología Tropical* 63 (Supplement 2): 37–47.
- Granja-Fernández R, Solís-Marín FA, Benítez-Villalobos F, Herrero-Pérezrul MD, López-Pérez A (2015b) Checklist of echinoderms (Echinodermata) from the Southern Mexican Pacific: a historical review. *Revista de Biología Tropical* 63 (Supplement 2): 87–114.
- Granja-Fernández R, Hendrickx ME, Rangel-Solís PD, López-Pérez A (2023) Deep-sea Ophiuroidea (Echinodermata) collected during the TALUD cruises in western Mexico. *Zootaxa* 5259: 1–71. <https://doi.org/10.11646/zootaxa.5259.1.1>
- Grupe BM, Krach ML, Pasulka AL, Maloney JM, Levin LA, Frieder CA (2015) Methane seep ecosystem functions and services from a recently discovered southern California seep. *Marine Ecology* 36: 91–108. <https://doi.org/10.1111/maec.12243>
- Gusmão LC, Daly M (2010) Evolution of sea anemones (Cnidaria: Actiniaria: Hormathiidae) symbiotic with hermit crabs. *Molecular Phylogenetics and Evolution* 56: 868–877. <https://doi.org/10.1016/j.ympev.2010.05.001>
- Gusmão LC, Van Deusen V, Daly M, Rodríguez E (2020) Origin and evolution of the symbiosis between sea anemones (Cnidaria, Anthozoa, Actiniaria) and hermit crabs, with additional notes on anemone-gastropod associations. *Molecular Phylogenetics and Evolution* 148: 106805. <https://doi.org/10.1016/j.ympev.2020.106805>
- Gustafson RG, Turner RD, Lutz RA, Vrijenhoek RC (1998) A new genus and five new species of mussels (Bivalvia, Mytilidae) from deep-sea sulfide/hydrocarbon seeps in the Gulf of Mexico. *Malacologia* 40: 63–112. <https://doi.org/10.1007/s11307-006-0060-5>
- Guzmán GL, Sellanes J (2011) *Spongicoloides* sp. aff. to *Spongicoloides galapagensis* (Decapoda: Stenopodidea: Spongicolidae): a new species for Chilean carcinofauna and the first record of a Stenopodid for the Chilean margin. *Latin American Journal of Aquatic Research* 39: 613–616. <https://doi.org/10.3856/vol39-issue3-fulltext-22>

- Guzman GL, Sellanes J (2015) A review of the Munidopsidae Ortmann, 1898 (Decapoda, Galatheoidea) in Chilean waters, including new records for the Southeastern Pacific. *Zootaxa* 4021: 282–306. <https://doi.org/10.11646/zootaxa.4021.2.3>
- Gwinn NE, Rinaldo C (2009) The Biodiversity Heritage Library: Sharing biodiversity literature with the world. *IFLA Journal* 35: 25–34. <https://doi.org/10.1177/0340035208102032>
- Han X, Suess E, Sahling H, Wallmann K (2004) Fluid venting activity on the Costa Rica margin: New results from authigenic carbonates. *International Journal of Earth Sciences* 93: 596–611. <https://doi.org/10.1007/s00531-004-0402-y>
- Handl C, Bouchet P (2007) Mystery tubes coiled around deep-water tropical gorgonians: Fecampiid cocoons (Platyhelminthes: Fecampiida) resembling Solenogastres (Mollusca). *Systematic Parasitology* 67: 81–85. <https://doi.org/10.1007/s11230-006-9077-z>
- Harry HW (1969) A review of the living leptonacean bivalves of the genus *Aligena*. *The Veliger* 11: 164–181. <https://www.biodiversitylibrary.org/page/42397993> [accessed on 27 May 2021]
- Hartman O (1955) Quantitative survey of the benthos of San Pedro Basin, southern California. Part I. Preliminary results. *Allan Hancock Pacific Expeditions* 19. <https://www.biodiversitylibrary.org/part/245138> [accessed on 25 April 2022]
- Hartman O (1961) Polychaetous annelids from California. *Allan Hancock Pacific Expeditions* 26: 1–226. <https://www.biodiversitylibrary.org/partpdf/19823> [accessed on 12 November 2023]
- Hartman O (1963) Submarine canyons of southern California. Part III. Systematics: Polychaetes. *Allan Hancock Pacific Expeditions* 27: 1–93. <https://www.biodiversitylibrary.org/page/4683344> [accessed on 29 March 2021]
- Hartman O, Barnard JL (1958) The benthic fauna of the deep basins off southern California. *Allan Hancock Pacific Expeditions* 22. <https://www.biodiversitylibrary.org/item/26532> [accessed on 24 April 2022]
- Hatch AS, Liew H, Hourdez S, Rouse GW (2020) Hungry scale worms: Phylogenetics of *Peinaleopolynoe* (Polynoidae, Annelida), with four new species. *ZooKeys* 932: 27–74. <https://doi.org/10.3897/zookeys.932.48532>
- Hayashi R (1939) Solasterids in Japanese waters. *Journal of the Faculty of Science of Hokkaido University* 6: 297–311. <http://hdl.handle.net/2115/27017> [accessed on 13 May 2021]
- He X, Xu T, Chen C, Liu X, Li Y-X, Zhong Z, Gu X, Lin Y-T, Lan Y, Yan G, Sun Y, Qiu J-W, Qian P-Y, Sun J (2023) Same (sea) bed different dreams: Biological community structure of the Haima seep reveals distinct biogeographic affinities. *The Innovation Geoscience* 1: 100019. <https://doi.org/10.59717/j.xinn-geo.2023.100019>
- Helgen LE, Rouse GW (2006) Species delimitation and distribution in *Aporometra* (Crinoidea : Echinodermata): Endemic Australian featherstars. *Invertebrate Systematics* 20: 395–414. <https://doi.org/10.1071/IS05050>
- Hemery LG, Roux M, Ameziane N, Eleaume M (2013) High-resolution crinoid phyletic inter-relationships derived from molecular data. *Cahiers de Biologie Marine* 54: 511–523.
- Henderson JR (1885) XXXIX.—Diagnoses of the new species of Galatheidea collected during the ‘Challenger’ expedition. *Annals and Magazine of Natural History* 16: 407–421. <https://doi.org/10.1080/00222938509459908>
- Hendrickx ME (1996) New records of deep-water decapod crustaceans in the southeastern Gulf of California, Mexico. *Revista de Biología Tropical* 44: 945–947. <https://revistas.ucr.ac.cr/index.php/rbt/article/view/32225> [accessed 3 May 2021]

- Hendrickx ME (2001) Occurrence of a continental slope decapod crustacean community along the edge of the minimum oxygen zone in the south eastern Gulf of California, Mexico. *Belgian Journal of Zoology* 131: 95–110. https://biblio.naturalsciences.be/associated_publications/bjz/131-2-supplement/volume-131-s2-pp-95-110.pdf/view [accessed on 3 May 2021]
- Hendrickx ME (Ed.) (2020a) *Deep-Sea Pycnogonids and Crustaceans of the Americas*. Springer, Cham, 708 pp. <https://doi.org/10.1007/978-3-030-58410-8>
- Hendrickx ME (2020b) The deep-water *Colossendeis tenera* Hilton, 1943 (Pycnogonida, Pantopoda, Colossendeidae) off western Mexico. In: Hendrickx ME (Ed.) *Deep-Sea Pycnogonids and Crustaceans of the Americas*. Springer, Cham, 39–45. https://doi.org/10.1007/978-3-030-58410-8_3
- Hendrickx ME, Ayon-Parente M (2009) Notes on distribution and ecology of *Parapagurus foraminosus* Lemaitre, 1999 (Decapoda, Anomura, Parapaguridae) in the eastern Pacific. *Crustaceana* 82: 1339–1344. <https://doi.org/10.1163/001121609X12481627024292>
- Hendrickx ME, Harvey AW (1999) Checklist of anomuran crabs (Crustacea: Decapoda) from the Eastern Tropical Pacific. *Belgian Journal of Zoology* 129: 363–389. https://biblio.naturalsciences.be/associated_publications/bjz/bibliographic-references/129-2/volume-129-2-pp-363-389.pdf [accessed on 18 June 2020]
- Hendrickx ME, Hernández-Payán JC (2018) Distribution and abundance of *Nematocarcinus* spp. (Crustacea: Decapoda: Caridea: Nematocarcinidae) off western Mexico, eastern Pacific. *Regional Studies in Marine Science* 23: 47–52. <https://doi.org/10.1016/j.rsma.2017.10.012>
- Hendrickx ME, López J (2020) Deep-water stomatopod and decapod crustaceans collected off Central America by the R/V “Miguel Oliver.” In: Hendrickx ME (Ed.) *Deep-Sea Pycnogonids and Crustaceans of the Americas*. Springer, Cham, 603–631. https://doi.org/10.1007/978-3-030-58410-8_23
- Hendrickx ME, Salgado-Barragán J (1991) Los estomatópodos (Crustacea: Hoplocarida) del Pacífico mexicano. *Publicaciones Especiales del Instituto de Ciencias del Mar y Limnología, UNAM* 10: 1–200.
- Hendrickx ME, Mah C, Zárate-Montes CM (2011) Deep-water Asteroidea (Echinodermata) collected during the TALUD cruises in the Gulf of California, Mexico. *Revista Mexicana de Biodiversidad* 82: 798–824. <https://doi.org/10.22201/ib.20078706e.2011.3.692>
- Hendrickx ME, Valentich-Scott P, Suárez-Mozo NY (2016) Deep-water bivalve mollusks collected during the TALUD XV cruise off the west coast of the southern Baja California Peninsula, Mexico. *Biodiversity Data Journal* 4: e8661. <https://doi.org/10.3897/BDJ.4.e8661>
- Hendrickx ME, Ayon-Parente M, Mejía-Mercado BE, Hinojosa-Corona A (2014) Registros de especies de crustáceos decápodos (Crustacea: Decapoda). In: Mejía-Mercado BE, Hinojosa-Corona A, Hendrickx ME (Eds), *Explorando el mar profundo del Golfo de California 2008-2014*. CICESE, Ensenada, Mexico, 176–190.
- Hernández P, Clarke TML, Benavides-Varela C, Villalobos-Rojas F, Nivia-Ruiz J, Wehrtmann IS (2011) Population demography and spatial distribution of the mantis shrimp *Squilla biformis* (Stomatopoda, Squillidae) from Pacific Costa Rica. *Marine Ecology Progress Series* 424: 157–168. <https://doi.org/10.3354/meps08982>
- Hertlein LG (1935) The Templeton Crocker Expedition of the California Academy of Sciences, 1932. No. 25. The Recent Pectinidae. *Proceedings of the California Academy of Sciences* 21: 301–328. <https://www.biodiversitylibrary.org/part/53429> [accessed on 28 May 2021]

- Herzig PM, Hanningen MD, Stoffers P, Becker K-P, Drischel M, Franklin J, Franz L, Gemmell JB, Hoppner B, Horn C, Horz K, Jellineck T, Jonasson IR, Kia P, Nickelsen S, Percival J, Perfit M, Petersen S, Schmidt M, Seifert T, Thiessen O, Turkay M, Tunnicliffe V, Winn K (1998) Petrology, gold mineralisation and biological communities at shallow submarine volcanoes of the New Ireland fore-arc (Papua-New Guinea): Preliminary results of R/V *Sonne* cruise SO-133. *InterRidge News* 7: 34–38. <https://oceanrep.geomar.de/id/eprint/33114/> [accessed on 12 March 2022]
- Hoareau TB, Boissin E (2010) Design of phylum-specific hybrid primers for DNA barcoding: Addressing the need for efficient COI amplification in the Echinodermata. *Molecular Ecology Resources* 10: 960–967. <https://doi.org/10.1111/j.1755-0998.2010.02848.x>
- Hooker Y, Elba P-R, Solís-Marín FA (2014) Echinoderms of Peru. In: Alvarado JJ, Solís-Marín FA (Eds), *Echinoderm Research and Diversity in Latin America*. Springer Berlin, Heidelberg, 277–299. https://doi.org/10.1007/978-3-642-20051-9_8
- Höpel CG, Yeo D, Grams M, Meier R, Richter S (2022) Mitogenomics supports the monophyly of Mysidacea and Peracarida (Malacostraca). *Zoologica Scripta*, 1–11. <https://doi.org/10.1111/zsc.12554>
- Horton T, Marsh L, Bett BJ, Gates AR, Jones DOB, Benoist NMA, Pfeifer S, Simon-Lledó E, Durden JM, Vandepitte L, Appeltans W (2021) Recommendations for the standardisation of open taxonomic nomenclature for image-based identifications. *Frontiers in Marine Science* 8: 620702. <https://doi.org/10.3389/fmars.2021.620702>
- Horton T, Lowry J, De Broyer C, Bellan-Santini D, Coleman CO, Corbari L, Costello MJ, Daneliya M, Dauvin J-C, Fišer C, Gasca R, Grabowski M, Guerra-García JM, Hendrycks E, Hughes L, Jaume D, Jazdzewski K, Kim Y-H, King R, Krapp-Schickel T, LeCroy S, Lörz A-N, Mamos T, Senna AR, Serejo C, Sket B, Souza-Filho JF, Tandberg AH, Thomas JD, Thurston M, Vader W, Väinölä R, Vonk R, White K, Zeidler W (2022) World Amphipoda Database. World Register of Marine Species. <https://www.marinespecies.org/aphia.php?p=taxdetails&id=1135> [accessed on 12 May 2022]
- Howell KL, Hilário A, Allcock AL, Bailey DM, Baker M, Clark MR, Colaço A, Copley J, Cordes EE, Danovaro R, Dissanayake A, Escobar E, Esquete P, Gallagher AJ, Gates AR, Gaudron SM, German CR, Gjerde KM, Higgs ND, Le Bris N, Levin LA, Manea E, McClain C, Menot L, Mestre NC, Metaxas A, Milligan RJ, Muthumbi Awn, Narayanaswamy BE, Ramalho SP, Ramirez-Llodra E, Robson LM, Rogers AD, Sellanes J, Sigwart JD, Sink K, Snelgrove PVR, Stefanoudis PV, Sumida PY, Taylor ML, Thurber AR, Vieira RP, Watanabe HK, Woodall LC, Xavier JR (2020) A blueprint for an inclusive, global deep-sea Ocean Decade field program. *Frontiers in Marine Science* 7: 584861. <https://doi.org/10.3389/fmars.2020.584861>
- Hugall AF, O'Hara TD, Hunjan S, Nilsen R, Moussalli A (2016) An exon-capture system for the entire class Ophiuroidea. *Molecular Biology and Evolution* 33: 281–294. <https://doi.org/10.1093/molbev/msv216>
- Irisarri I, Uribe JE, Eernisse DJ, Zardoya R (2020) A mitogenomic phylogeny of chitons (Mollusca: Polyplacophora). *BMC Evolutionary Biology* 20: 1–15. <https://doi.org/10.1186/s12862-019-1573-2>
- Jarms G, Morandini AC, da Silveira FL (2002) Polyps of the families Atorellidae and Nausithoidae (Scyphozoa: Coronatae) new to the Brazilian fauna. *Biota Neotropica* 2: 1–11. <https://doi.org/10.1590/S1676-06032002000100004>
- Jereb P, Roper CFE (2010) *Cephalopods of the world. An annotated and illustrated catalogue of cephalopod species known to date. Volume 2. Myopsid and oegopsid squids*. FAO, Rome. <https://www.fao.org/publications/card/en/c/42d0f8a9-1696-5dcb-a323-e8303399a472/> [accessed on 28 March 2022]

- Johnson HP (1901) The Polychaeta of the Puget Sound region. Proceedings of the Boston Society of Natural History 29: 381–437. <https://www.marinespecies.org/aphia.php?p=sourcedetails&id=50546> [accessed on 5 May 2020]
- Johnson SB, Young CR, Jones WJ, Warén A, Vrijenhoek RC (2006) Migration, isolation, and speciation of hydrothermal vent limpets (Gastropoda; Lepetodrilidae) across the Blanco Transform Fault. The Biological Bulletin 210: 140–157. <https://doi.org/10.2307/4134603>
- Johnson SB, Warén A, Vrijenhoek RC (2008) DNA barcoding of *Lepetodrilus* limpets reveals cryptic species. Journal of Shellfish Research 27: 43–51. [https://doi.org/10.2983/0730-8000\(2008\)27\[43:DBOLLR\]2.0.CO;2](https://doi.org/10.2983/0730-8000(2008)27[43:DBOLLR]2.0.CO;2)
- Johnson SB, Won YJ, Harvey JB, Vrijenhoek RC (2013) A hybrid zone between *Bathymodiolus* mussel lineages from eastern Pacific hydrothermal vents. BMC Evolutionary Biology 13: 21. <https://doi.org/10.1186/1471-2148-13-21>
- Johnson SB, Krylova EM, Audzijonyte A, Sahling H, Vrijenhoek RC (2017) Phylogeny and origins of chemosynthetic vesicomyid clams. Systematics and Biodiversity 15: 346–360. <https://doi.org/10.1080/14772000.2016.1252438>
- Jones WJ, Macpherson E (2007) Molecular phylogeny of the East Pacific squat lobsters of the genus *Munidopsis* (Decapoda: Galatheididae) with the descriptions of seven new species. Journal of Crustacean Biology 27: 477–501. <https://doi.org/10.1651/S-2791.1>
- Kahn LM, Silver EA, Orange D, Kochevar R, McAdoo B (1996) Surficial evidence of fluid expulsion from the Costa Rica accretionary prism. Geophysical Research Letters 23: 887–890. <https://doi.org/10.1029/96GL00732>
- Kamenev GM (2013) Species composition and distribution of bivalves in bathyal and abyssal depths of the Sea of Japan. Deep Sea Research Part II: Topical Studies in Oceanography 86–87: 124–139. <https://doi.org/10.1016/j.dsr2.2012.08.004>
- Karaseva NP, Rimskaya-Korsakova NN, Galkin SV, Malakhov VV (2016) Taxonomy, geographical and bathymetric distribution of vestimentiferan tubeworms (Annelida, Siboglinidae). Biology Bulletin 43: 937–969. <https://doi.org/10.1134/S1062359016090132>
- Katoh K, Rozewicki J, Yamada KD (2018) MAFFT online service: Multiple sequence alignment, interactive sequence choice and visualization. Briefings in Bioinformatics 20: 1160–1166. <https://doi.org/10.1093/bib/bbx108>
- Kenk VC, Wilson BR (1985) A new mussel (Bivalvia, Mytilidae) from hydrothermal vents in the Galapagos Rift zone. Malacologia 26: 253–271. <https://www.biodiversitylibrary.org/page/13135080> [accessed on 29 May 2020]
- Kiel S (2016) A biogeographic network reveals evolutionary links between deep-sea hydrothermal vent and methane seep faunas. Proceedings of the Royal Society B: Biological Sciences 283: 20162337. <https://doi.org/10.1098/rspb.2016.2337>
- Kimura G, Silver E, Blum P, Blanc G, Bolton A, Clennell M, Griffin J, Housen B, Ibaraki M, Kanamatsu T, Kastner M, Lindsley-Griffin N, Lueckge A, McIntosh K, Meschede M, Morris J, Muza J, Myers G, Protti M, Saether O, Saito S, Scholl D, Spence G, Tobin H, Vannucchi P, White L (1997) Introduction. In: Proceedings of the Ocean Drilling Program, 170 Initial Reports. Ocean Drilling Program, 7–17. <https://doi.org/10.2973/odp.proc.ir.170.101.1997>
- Knott KE, Wray GA (2000) Controversy and consensus in asteroid systematics: New insights to ordinal and familial relationships. American Zoologist 40: 382–392. <https://doi.org/10.1093/icb/40.3.382>

- Kobayashi G, Araya JF (2018) Southernmost records of *Escarpia spicata* and *Lamellibrachia barhami* (Annelida: Siboglinidae) confirmed with DNA obtained from dried tubes collected from undiscovered reducing environments in northern Chile. PLoS ONE 13: e0204959. <https://doi.org/10.1371/journal.pone.0204959>
- Kocot KM, Todt C, Mikkelsen NT, Halanych KM (2019) Phylogenomics of Aplacophora (Mollusca, Aculifera) and a solenogaster without a foot. Proceedings of the Royal Society B: Biological Sciences 286: 20190115. <https://doi.org/10.1098/rspb.2019.0115>
- Koga C, Rouse GW (2021) Mitogenomics and the phylogeny of mantis shrimps (Crustacea: Stomatopoda). Diversity 13: 647. <https://doi.org/10.3390/d13120647>
- Kongsrud JA, Rapp HT (2012) *Nicomache (Loxochona) lokii* sp. nov. (Annelida: Polychaeta: Maldanidae) from the Loki's Castle vent field: An important structure builder in an Arctic vent system. Polar Biology 35: 161–170. <https://doi.org/10.1007/s00300-011-1048-4>
- Kou Q, Meland K, Li X, He L, Wang Y (2019) Deepest record of *Eucopia sculpticauda* (Crustacea: Lophogastrida: Eucopiidae) and the order, with new insights into the distribution and genetic diversity of the species. Bulletin of Marine Science 95: 327–335. <https://doi.org/10.5343/bms.2018.0074>
- Kroupa A, Remsen D (2010) Chapter 4 Individual records and the associated data: information standards and protocols. In: Eymann J, Degreef J, Häuser C, Monje JC, Samyn Y, Vandenspiegel D (Eds), Manual on Field Recording Techniques and Protocols for All Taxa Biodiversity Inventories. Abc Taxa 8: 49–67. <https://abctaxa.naturalsciences.be/pdf-8/> [accessed on 16 June 2021]
- Krylova EM, Sahling H (2006) Recent bivalve molluscs of the genus *Calyptogena* (Vesicomidae). The Journal of Molluscan Studies 72: 359–395. <https://doi.org/10.1093/mollus/eyl022>
- Kunze T (2011) *Dillwynella voightae* new species, a new skeneimorph gastropod (Turbinidae) from the western Atlantic and a new record of *Dillwynella modesta* (Dali, 1889). The Nautilus 125: 36–40. <https://www.biodiversitylibrary.org/part/174670> [accessed on 17 July 2020]
- Langeneck J, Barbieri M, Maltagliati F, Castelli A (2019) Molecular phylogeny of Paraeonidae (Annelida). Molecular Phylogenetics and Evolution 136: 1–13. <https://doi.org/10.1016/j.ympev.2019.03.023>
- Law CJ, Dorgan KM, Rouse GW (2014) Relating divergence in polychaete musculature to different burrowing behaviors: A study using Opheliidae (Annelida). Journal of Morphology 275: 548–571. <https://doi.org/10.1002/jmor.20237>
- Leigh JW, Bryant D (2015) POPART: Full-feature software for haplotype network construction. Methods in Ecology and Evolution 6: 1110–1116. <https://doi.org/10.1111/2041-210X.12410>
- Lemaitre R (2014) A worldwide taxonomic and distributional synthesis of the genus *Oncopagurus* Lemaitre, 1996 (Crustacea: Decapoda: Anomura: Parapaguridae), with descriptions of nine new species. The Raffles Bulletin of Zoology 62: 210–301. <https://repository.si.edu/handle/10088/21941> [accessed on 20 February 2024]
- Lessios HA (2005) Echinoids of the Pacific waters of Panama: status of knowledge and new records. Revista de Biología Tropical 53: 147–170.
- Levin LA, Rouse GW (2020) Giant protists (xenophyophores) function as fish nurseries. Ecology 101: e02933. <https://doi.org/10.1002/ecy.2933>
- Levin LA, James DW, Martin CM, Rathburn AE, Harris LH, Michener RH (2000) Do methane seeps support distinct macrofaunal assemblages? Observations on community

- structure and nutrition from the northern California slope and shelf. *Marine Ecology Progress Series* 208: 21–39. <https://doi.org/10.3354/meps208021>
- Levin LA, Mendoza GF, Konotchick T, Lee R (2009) Macrobenthos community structure and trophic relationships within active and inactive Pacific hydrothermal sediments. *Deep Sea Research Part II: Topical Studies in Oceanography* 56: 1632–1648. <https://doi.org/10.1016/j.dsr2.2009.05.010>
- Levin LA, Orphan VJ, Rouse GW, Rathburn AE, Ussler W, Cook GS, Goffredi SK, Perez EM, Warén A, Grupe BM, Chadwick G, Strickrott B (2012) A hydrothermal seep on the Costa Rica margin: Middle ground in a continuum of reducing ecosystems. *Proceedings of the Royal Society B: Biological Sciences* 279: 2580–2588. <https://doi.org/10.1098/rspb.2012.0205>
- Levin LA, Mendoza GF, Grupe BM, Gonzalez JP, Jellison B, Rouse G, Thurber AR, Warén A (2015) Biodiversity on the rocks: Macrofauna inhabiting authigenic carbonate at Costa Rica methane seeps. *PLoS ONE* 10: e0131080. <https://doi.org/10.1371/journal.pone.0131080>
- Levin LA, Baco AR, Bowden DA, Colaco A, Cordes EE, Cunha MR, Demopoulos AWJ, Gobin J, Grupe BM, Le J, Metaxas A, Netburn AN, Rouse GW, Thurber AR, Tunnicliffe V, Van Dover CL, Vanreusel A, Watling L (2016) Hydrothermal vents and methane seeps: Rethinking the sphere of influence. *Frontiers in Marine Science* 3: 72. <https://doi.org/10.3389/fmars.2016.00072>
- Levin LA, Mendoza GF, Grupe BM (2017) Methane seepage effects on biodiversity and biological traits of macrofauna inhabiting authigenic carbonates. *Deep Sea Research Part II: Topical Studies in Oceanography* 137: 26–41. <https://doi.org/10.1016/j.dsr2.2016.05.021>
- Liang J, Feng JC, Zhang S, Cai Y, Yang Z, Ni T, Yang HY (2021) Role of deep-sea equipment in promoting the forefront of studies on life in extreme environments. *iScience* 24: 1–21. <https://doi.org/10.1016/j.isci.2021.103299>
- Linchangco GV, Foltz DW, Reid R, Williams J, Nodzak C, Kerr AM, Miller AK, Hunter R, Wilson NG, Nielsen WJ, Mah CL, Rouse GW, Wray GA, Janies DA (2017) The phylogeny of extant starfish (Asteroidea: Echinodermata) including *Xyloplax*, based on comparative transcriptomics. *Molecular Phylogenetics and Evolution* 115: 161–170. <https://doi.org/10.1016/j.ympev.2017.07.022>
- Lindgren J, Hatch AS, Hourdez S, Seid CA, Rouse GW (2019) Phylogeny and biogeography of *Branchipolynoe* (Polynoidae, Phyllococida, Aciculata, Annelida), with descriptions of five new species from methane seeps and hydrothermal vents. *Diversity* 11(9): 153. <https://doi.org/10.3390/d11090153>
- Linse K, Jackson JA, Fitzcharles E, Sands CJ, Buckeridge JS (2013) Phylogenetic position of Antarctic Scalpelliformes (Crustacea: Cirripedia: Thoracica). *Deep Sea Research Part I: Oceanographic Research Papers* 73: 99–116. <https://doi.org/10.1016/j.dsr.2012.11.006>
- Liu H, Cai S, Liu J, Zhang H (2018) Comparative mitochondrial genomic analyses of three chemosynthetic vesicomyid clams from deep-sea habitats. *Ecology and Evolution* 8: 7261–7272. <https://doi.org/10.1002/ece3.4153>
- Lopes DA, Hajdu E, Reiswig HM (2011) Taxonomy of *Farrea* (Porifera, Hexactinellida, Hexactinosida) from the southwestern Atlantic, with description of a new species and a discussion on the recognition of subspecies in Porifera. *Canadian Journal of Zoology* 89: 169–189. <https://doi.org/10.1139/Z10-105>
- Lorion J, Kiel S, Faure B, Kawato M, Ho SYW, Marshall B, Tsuchida S, Miyazaki JI, Fujiwara Y (2013) Adaptive radiation of chemosymbiotic deep-sea mussels. *Proceedings*

- of the Royal Society B: Biological Sciences 280: 20131243. <https://doi.org/10.1098/rspb.2013.1243>
- Ludwig H (1893) Vorläufiger Bericht über die erbeuteten Holothurien. In No. 4: Reports on the dredging operations off the west coast of Central America to the Galapagos, to the west coast of Mexico, and in the Gulf of California, in charge of Alexander Agassiz, carried on by the U.S. Fish Commission steamer "Albatross," during 1891, Lieut. Commander Z. L. Tanner, U. S. N., Commanding. Bulletin of the Museum of Comparative Zoology at Harvard College 24: 105–114. <https://www.biodiversitylibrary.org/page/28865930> [accessed on 13 April 2021]
- Ludwig H (1894) Reports on an exploration off the west coasts of Mexico, Central and South America, and off the Galapagos Islands, in charge of Alexander Agassiz, by the U.S. Fish Commission steamer "Albatross," during 1891, Lt. Commander Z.L. Tanner, U.S.N., commanding. XII. The Holothuriodea. Memoirs of the Museum of Comparative Zoölogy at Harvard College 17: 183. <https://www.biodiversitylibrary.org/page/4339195> [accessed on 23 May 2020]
- Ludwig H (1905) Reports on the exploration off the west coasts of Mexico, Central and South America, and off the Galapagos Islands, in charge of Alexander Agassiz by the U.S. Fish Commission Steamer "Albatross," during 1891, Lieut.-Commander Z.L. Tanner, U.S.N., Commanding. XXXV. Reports on the scientific results of the expedition to the tropical Pacific in charge of Alexander Agassiz, by the U.S. Fish Commission Steamer "Albatross", from August, 1899, to March, 1900, Commander Jefferson F. Moser, U.S. N., Commanding. VII. Asteroidea. Memoirs of the Museum of Comparative Zoology at Harvard College 32: 1–292. <https://doi.org/10.5962/bhl.part.27494>
- Lundsten L, Schlining KL, Frasier K, Johnson SB, Kuhn LA, Harvey JBJ, Clague G, Vrijenhoek RC (2010) Time-series analysis of six whale-fall communities in Monterey Canyon, California, USA. Deep Sea Research Part I: Oceanographic Research Papers 57: 1573–1584. <https://doi.org/10.1016/j.dsr.2010.09.003>
- Lutken CF, Mortensen T (1899) Reports on an exploration off the west coasts of Mexico, central and South America, and the Galapagos Islands, in charge of Alexander Agassiz, by the U.S. Fish Commission steamer "Albatross," during 1891, Lieut. Commander Z.L. Tanner, U.S.N., commanding. XXV. The Ophiuridae. Memoirs of the Museum of Comparative Zoology 23: 97–208. <https://www.biodiversitylibrary.org/page/28891692> [accessed on 27 May 2021]
- Lyman T (1878) Ophiuridae and Astrophytidae of the exploring voyage of H. M. S. "Challenger," under Prof. Sir Wyville Thomson, F. R. S. Part I. Bulletin of the Museum of Comparative Zoology at Harvard College 5: 65–168. <https://www.biodiversitylibrary.org/page/28876644> [accessed on 27 May 2021]
- Lyman T (1879) Ophiuridae and Astrophytidae of the exploring voyage of H.M.S. "Challenger," under Prof. Sir Wyville Thomson, F.R.S. Part II. Bulletin of the Museum of Comparative Zoology at Harvard College 6: 17–83. <https://www.biodiversitylibrary.org/page/31068682> [accessed on 20 April 2022]
- MacPherson E, Wehrmann IS (2010) Occurrence of lithodid crabs (Decapoda, Lithodidae) on the Pacific coast of Costa Rica, Central America. Crustaceana 83: 143–151. <https://doi.org/10.1163/001121609X12487811051660>
- Mah CL (2016) Deep-sea (>1000 m) Goniasteridae (Valvatida; Asteroidea) from the North Pacific, including an overview of *Sibogaster*, *Bathyceramaster* n. gen. and three new species. Zootaxa 4175: 104–141. <https://doi.org/10.11646/zootaxa.4175.2.1>
- Mah C, Nizinski M, Lundsten L (2010) Phylogenetic revision of the Hippasterinae (Goniasteridae; Asteroidea): Systematics of deep sea corallivores, including one new

- genus and three new species. *Zoological Journal of the Linnean Society* 160: 266–301. <https://doi.org/10.1111/j.1096-3642.2010.00638.x>
- Marlow JJ, Anderson RE, Reysenbach AL, Seewald JS, Shank TM, Teske AP, Wanless VD, Soule SA (2022) New opportunities and untapped scientific potential in the abyssal ocean. *Frontiers in Marine Science* 8: 798943. <https://doi.org/10.3389/fmars.2021.798943>
- Martin AM, Goffredi SK (2012) “*Pliocardia*” *krylovata*, a new species of vesicomid clam from cold seeps along the Costa Rica Margin. *Journal of the Marine Biological Association of the United Kingdom* 92: 1127–1137. <https://doi.org/10.1017/S0025315411000713>
- Martin JW, Wall AR, Shank T, Cha H, Seid CA, Rouse GW (2018) A new species of *Alvinocaris* (Crustacea: Decapoda: Caridea: Alvinocarididae) from Costa Rican methane seeps. *Zootaxa* 4504: 418–430. <https://doi.org/10.11646/zootaxa.4504.3.7>
- Martínez-Guerrero B, López-Pérez A (2018) Richness and large-scale distribution of marine benthic caridean shrimps (Decapoda: Caridea) from the Eastern Tropical Pacific. *Nauplius* 26: 1–16. <https://doi.org/10.1590/2358-2936e2018035>
- Martins L, Tavares M (2018) *Ypsilothuria bitentaculata bitentaculata* (Echinodermata: Holothuroidea) from the southwestern Atlantic, with comments on its morphology. *Zoologia* 35: e24573. <https://doi.org/10.3897/zoologia.35.e24573>
- Massin C, Hendrickx ME (2011) Deep-water Holothuroidea (Echinodermata) collected during the TALUD cruises off the Pacific coast of Mexico, with the description of two new species. *Revista Mexicana de Biodiversidad* 82: 413–443. <https://doi.org/10.22201/ib.20078706e.2011.2.476>
- Matabos M, Jollivet D (2019) Revisiting the *Lepetodrilus elevatus* species complex (Vetigastropoda: Lepetodrilidae), using samples from the Galápagos and Guaymas hydrothermal vent systems. *Journal of Molluscan Studies* 85: 66–72. <https://doi.org/10.1093/mollus/eyy061>
- Mau S, Sahling H, Rehder G, Suess E, Linke P, Soeding E (2006) Estimates of methane output from mud extrusions at the erosive convergent margin off Costa Rica. *Marine Geology* 225: 129–144. <https://doi.org/10.1016/j.margeo.2005.09.007>
- Mau S, Rehder G, Sahling H, Schleicher T, Linke P (2014) Seepage of methane at Jaco Scar, a slide caused by seamount subduction offshore Costa Rica. *International Journal of Earth Sciences* 103: 1801–1815. <https://doi.org/10.1007/s00531-012-0822-z>
- Maxwell J, Taboada S, Taylor ML (2022) *Gorgoniapolynoe caeciliae* revisited: The discovery of new species and molecular connectivity in deep-sea commensal polynoids from the Central Atlantic. *Deep Sea Research Part I: Oceanographic Research Papers* 185: 103804. <https://doi.org/10.1016/j.dsr.2022.103804>
- McAdoo BG, Orange DL, Silver EA, McIntosh K, Abbott L, Galewsky J, Kahn L, Protti M (1996) Seafloor structural observations, Costa Rica Accretionary Prism. *Geophysical Research Letters* 23: 883–886. <https://doi.org/10.1029/96GL00731>
- McClain C, Nekola J (2007) Notes on two cases of molluscan biogeography. *American Conchologist* 35: 4–7. <https://www.biodiversitylibrary.org/page/62541708> [accessed on 7 July 2024]
- McClain CR, Boolukos CM, Bryant SRD, Hanks G (2023) Sunken trees in the deep sea link terrestrial and marine biodiversity. *Ecology* 104: e4168. <https://doi.org/10.1002/ecy.4168>
- McCowin MF, Rouse GW (2018) A new *Lamellibrachia* species and confirmed range extension for *Lamellibrachia barhami* (Siboglinidae, Annelida) from Costa Rica methane seeps. *Zootaxa* 4504: 1–22. <https://doi.org/10.11646/zootaxa.4504.1.1>

- McCowin MF, Feehery C, Rouse GW (2020) Spanning the depths or depth-restricted: Three new species of *Bathymodiolus* (Bivalvia, Mytilidae) and a new record for the hydrothermal vent *Bathymodiolus thermophilus* at methane seeps along the Costa Rica margin. Deep Sea Research Part I: Oceanographic Research Papers 164: 103322. <https://doi.org/10.1016/j.dsr.2020.103322>
- McFadden CS, Quattrini AM, Brugler MR, Cowman PF, Dueñas LF, Kitahara MV, Paz-García DA, Reimer JD, Rodríguez E (2021) Phylogenomics, origin, and diversification of anthozoans (Phylum Cnidaria). Systematic Biology 70: 635–647. <https://doi.org/10.1093/sysbio/syaa103>
- McFadden CS, van Ofwegen LP, Quattrini AM (2022) Revisionary systematics of Octocorallia (Cnidaria: Anthozoa) guided by phylogenomics. Bulletin of the Society of Systematic Biologists 1: 8735. <https://doi.org/10.18061/bssb.v1i3.8735>
- McLean JH (1988) New archaeogastropod limpets from hydrothermal vents; superfamily Lepetodrilacea I. Systematic descriptions. Philosophical Transactions of the Royal Society of London. Series B, Biological Sciences 319: 1–32. <https://doi.org/10.1098/rstb.1988.0031>
- McLean JH (1992) Cocculiniform limpets (Cocculinidae and Pyropeltidae) living on whale bone in the deep sea off California. Journal of Molluscan Studies 58: 401–415. <https://doi.org/10.1093/mollus/58.4.401>
- McLean JH (1993) New species and records of *Lepetodrilus* (Vetigastropoda: Lepetodrilidae) from hydrothermal vents. The Veliger 36: 27–35. <https://www.biodiversitylibrary.org/page/42465631> [accessed on 11 June 2020]
- McLean JH (1996) The Prosobranchia. In: Blake JA, Scott PH, Lissner AL (Eds), Taxonomic Atlas of the Benthic Fauna of the Santa Maria Basin and Western Santa Barbara Channel. Volume 9. The Mollusca Part 2 – The Gastropoda. Santa Barbara Museum of Natural History, CA, USA, 1-160.
- McLean JH (2008) Three new species of the family Neolepetopsidae (Patellogastropoda) from hydrothermal vents and whale falls in the northeastern Pacific. Journal of Shellfish Research 27: 15–20. [https://doi.org/10.2983/0730-8000\(2008\)27\[15:TNSOTF\]2.0.CO;2](https://doi.org/10.2983/0730-8000(2008)27[15:TNSOTF]2.0.CO;2)
- McLean JH, Haszprunar G (1987) Pyropeltidae, a new family of cocculiniform limpets from hydrothermal vents. The Veliger 30: 196–205. <https://www.biodiversitylibrary.org/part/94179> [accessed on 22 June 2020]
- McMullin ER, Hourdez S, Schaeffer SW, Fisher CR (2003) Phylogeny and biogeography of deep sea vestimentiferan tubeworms and their bacterial symbionts. Symbiosis 34: 1–41. <http://hdl.handle.net/10222/77925> [accessed on 4 May 2020]
- Messing CG, White CM (2001) A revision of the Zenometridae (new rank) (Echinodermata, Crinoidea, Comatulidina). Zoologica Scripta 30: 159–180. <https://doi.org/10.1046/j.1463-6409.2001.00062.x>
- Metcalf KS, Murali R, Mullin SW, Connon SA, Orphan VJ (2021) Experimentally-validated correlation analysis reveals new anaerobic methane oxidation partnerships with consortium-level heterogeneity in diazotrophy. The ISME Journal 15: 377–396. <https://doi.org/10.1038/s41396-020-00757-1>
- Meyer CP (2003) Molecular systematics of cowries (Gastropoda: Cypraeidae) and diversification patterns in the tropics. Biological Journal of the Linnean Society 79: 401–459. <https://doi.org/10.1046/j.1095-8312.2003.00197.x>
- Mikkelsen NT, Todt C, Kocot KM, Halanych KM, Willassen E (2019) Molecular phylogeny of Caudofoveata (Mollusca) challenges traditional views. Molecular Phylogenetics and Evolution 132: 138–150. <https://doi.org/10.1016/j.ympev.2018.10.037>

- Miller A, Virmani JI (2023) Advanced marine technologies for ocean research. Deep Sea Research Part II: Topical Studies in Oceanography 212: 105340. <https://doi.org/10.1016/j.dsr2.2023.105340>
- Miller AK, Kerr AM, Paulay G, Reich M, Wilson NG, Carvajal JI, Rouse GW (2017) Molecular phylogeny of extant Holothuroidea (Echinodermata). Molecular Phylogenetics and Evolution 111: 110–131. <https://doi.org/10.1016/j.ympev.2017.02.014>
- Mills S, Mackay K (2019) *Caenopedina hawaiiensis* occurrences. NIWA Invertebrate Collection. Version 1.2. The National Institute of Water and Atmospheric Research (NIWA). Gbif.org occurrence dataset. <https://doi.org/10.15468/6qgswo>
- Milne Edwards H (1837) Histoire naturelle des crustacés, comprenant l'anatomie, la physiologie et la classification de ces animaux. Encyclopédique Roret 3: 638. <https://doi.org/10.5962/bhl.title.6234>
- Miloslavich P, Klein E, Díaz JM, Hernández CE, Bigatti G, Campos L, Artigas F, Castillo J, Penchaszadeh PE, Neill PE, Carranza A, Retana MV, Díaz de Astarloa JM, Lewis M, Yorio P, Piriz ML, Rodríguez D, Valentin YY, Gamboa L, Martín A (2011) Marine biodiversity in the Atlantic and Pacific coasts of South America: Knowledge and gaps. PLoS ONE 6(1): e14631. <https://doi.org/10.1371/journal.pone.0014631>
- Miura T, Ohta S (1991) Two polychaete species from the deep-sea hydrothermal vent in the middle Okinawa Trough. Zoological Science 8: 383–387. <https://www.biodiversitylibrary.org/part/71628> [accessed on 4 May 2020]
- Molinari CG, Collins AG, Morandini AC (2023) A morphological review of the jellyfish genus *Nausithoe* Kölliker, 1853 (Nausithoideae, Coronatae, Scyphozoa, Cnidaria). Zootaxa 5336: 1–32. <https://doi.org/10.11646/zootaxa.5336.1.1>
- Mongiardino Koch N, Coppard SE, Lessios HA, Briggs DEG, Mooi R, Rouse GW (2018) A phylogenomic resolution of the sea urchin tree of life. BMC Evolutionary Biology 18: 189. <https://doi.org/10.1186/s12862-018-1300-4>
- Mongiardino Koch N, Tilic E, Miller AK, Stiller J, Rouse GW (2023) Confusion will be my epitaph: genome-scale discordance stifles phylogenetic resolution of Holothuroidea. Proceedings of the Royal Society B: Biological Sciences 290: 20230988. <https://doi.org/10.1098/rspb.2023.0988>
- Mooi R, Constable H, Lockhart S, Pearse J (2004) Echinothurioid phylogeny and the phylogenetic significance of Kamptosoma (Echinoidea: Echinodermata). Deep Sea Research Part II: Topical Studies in Oceanography 51: 1903–1919. <https://doi.org/10.1016/j.dsr2.2004.07.020>
- Moore JP (1923) The polychaetous annelids dredged by the U.S.S. "Albatross" off the coast of southern California in 1904. Proceedings of the Academy of Natural Sciences of Philadelphia 75. <https://doi.org/10.5962/bhl.title.12425>
- Moore JM, Nishi E, Rouse GW (2017) Phylogenetic analyses of Chaetopteridae (Annelida). Zoologica Scripta 46: 596–610. <https://doi.org/10.1111/zsc.12238>
- Munilla T, Soler Membrives A (2009) Check-list of the pycnogonids from Antarctic and sub-Antarctic waters: zoogeographic implications. Antarctic Science 21: 99–111. <https://doi.org/10.1017/S095410200800151X>
- Murray J, Thomson CW (1885) Narrative Volume 1. Second Part. Appendix II. In: Report on the scientific results of the voyage of H. M. S. Challenger during the years 1873-76 under the command of Captain George S. Nares, R.N., F.R.S. and the late Captain Frank Tourle Thomson, R. N. Neill, Edinburgh, 1007–1015. <https://doi.org/10.5962/bhl.title.60036>
- National Academies of Sciences Engineering and Medicine (2020) Biological Collections: Ensuring Critical Research and Education for the 21st Century. National Academies Press, Washington, D.C. <https://doi.org/10.17226/25592>

- National Science and Technology Council Committee on Science Interagency Working Group on Scientific Collections (2009) Scientific collections: Mission-critical infrastructure for federal science agencies. Office of Science and Technology Policy, Washington, D.C. https://iwgsc.nal.usda.gov/sites/default/files/IWGSC_GreenReport_FINAL_2009.pdf [accessed on 15 November 2017]
- Neira C, Ingels J, Mendoza G, Hernandez-Lopez E, Levin LA (2018) Distribution of meiofauna in bathyal sediments influenced by the oxygen minimum zone off Costa Rica. *Frontiers in Marine Science* 5: 448. <https://doi.org/10.3389/fmars.2018.00448>
- Neulinger SC, Sahling H, Süling J, Imhoff JF (2006) Presence of two phylogenetically distinct groups in the deep-sea mussel *Acharax* (Mollusca: Bivalvia: Solemyidae). *Marine Ecology Progress Series* 312: 161–168. <https://doi.org/10.3354/meps312161>
- Niemann H, Linke P, Knittel K, MacPherson E, Boetius A, Brückmann W, Larvik G, Wallmann K, Schacht U, Omeregge E, Hilton D, Brown K, Rehder G (2013) Methane-carbon flow into the benthic food web at cold seeps – a case study from the Costa Rica subduction zone. *PLoS ONE* 8: e74894. <https://doi.org/10.1371/journal.pone.0074894>
- Norén M, Jondelius U (1999) Phylogeny of the Prolecithophora (Platyhelminthes) inferred from 18S rDNA sequences. *Cladistics* 15: 103–112. <https://doi.org/10.1111/j.1096-0031.1999.tb00252.x>
- O’Dea A, Lessios HA, Coates AG, Eytan RI, Restrepo-Moreno SA, Cione AL, Collins LS, De Queiroz A, Farris DW, Norris RD, Stallard RF, Woodburne MO, Aguilera O, Aubry MP, Berggren WA, Budd AF, Cozzuol MA, Coppard SE, Duque-Caro H, Finnegan S, Gasparini GM, Grossman EL, Johnson KG, Keigwin LD, Knowlton N, Leigh EG, Leonard-Pingel JS, Marko PB, Pyenson ND, Rachello-Dolmen PG, Soibelzon E, Soibelzon L, Todd JA, Vermeij GJ, Jackson JBC (2016) Formation of the Isthmus of Panama. *Science Advances* 2: e1600883. <https://doi.org/10.1126/sciadv.1600883>
- O’Hara TD, Hugall AF, Thuy B, Stöhr S, Martynov AV (2017) Restructuring higher taxonomy using broad-scale phylogenomics: The living Ophiuroidea. *Molecular Phylogenetics and Evolution* 107: 415–430. <https://doi.org/10.1016/j.ympev.2016.12.006>
- O’Hara TD, Stöhr S, Hugall AF, Thuy B, Martynov A (2018) Morphological diagnoses of higher taxa in Ophiuroidea (Echinodermata) in support of a new classification. *European Journal of Taxonomy* 416: 1–35. <https://doi.org/10.5852/ejt.2018.416>
- O’Loughlin PM, Paulay G, Davey N, Michonneau F (2011) The Antarctic region as a marine biodiversity hotspot for echinoderms: Diversity and diversification of sea cucumbers. *Deep Sea Research Part II: Topical Studies in Oceanography* 58: 264–275. <https://doi.org/10.1016/j.dsr2.2010.10.011>
- Ogawa A, Kremenetskaia A, Hiruta SF, Shibata Y, Narimatsu Y, Miki S, Morita T, Tsuchida S, Fujiwara Y, Fujita T (2022) Rehabilitation of two deep-sea holothurian species in genus *Pannychia* from the northwest Pacific Ocean. *Deep Sea Research Part II: Topical Studies in Oceanography* 202: 105099. <https://doi.org/10.1016/j.dsr2.2022.105099>
- Ogawa A, Kobayashi I, Kohtsuka H, Fujita T (2023) Two new species of the bathyal holothurian genus *Pannychia* (Elasipodida, Laetmogonidae) from Japanese waters. *Zootaxa* 5323: 105–125. <https://doi.org/10.11646/zootaxa.5323.1.6>
- Okutani T, Koshi-ishi T, Sato T, Imai T, Kato C (2009) Vesicomid fauna in the Chishima (Kurile) Trench: Occurrences of a new taxon and *Calyptogena extenta*. *Venus* 68: 15–25. https://doi.org/10.18941/venus.68.1-2_15
- Oliver PG, Sellanes J (2005) New species of Thyasiridae from a methane seepage area off Concepción, Chile. *Zootaxa* 1092: 1–20. <https://doi.org/10.11646/zootaxa.1092.1.1>
- Olu K, Duperret A, Sibuet M, Foucher JP, Fiala-Médioni A (1996) Structure and distribution of cold seep communities along the Peruvian active margin: Relationship to

- geological and fluid patterns. *Marine Ecology Progress Series* 132: 109–125. <https://doi.org/10.3354/meps132109>
- Opresko DM, Breedy O (2010) A new species of antipatharian coral (Cnidaria: Anthozoa: Antipatharia: Schizopathidae) from the Pacific coast of Costa Rica. *Proceedings of the Biological Society of Washington* 123: 234–241. <https://doi.org/10.2988/10-06.1>
- Orphan VJ, House CH, Hinrichs K-U, McKeegan KD, DeLong EF (2001) Methane-consuming archaea revealed by directly coupled isotopic and phylogenetic analysis. *Science* 293: 484–487. <https://doi.org/10.1126/science.1061338>
- Palumbi SR (1996) Nucleic Acids II: The Polymerase Chain Reaction. In: Hillis DM, Moritz C, Mable BK (Eds), *Molecular Systematics*, 2nd edn. Sinauer Associates, Sunderland, Massachusetts, 205–246. [accessed on 15 June 2021]
- Palumbi S, Martin A, Romano S, Mcmillan WO, Stice L, Grabowski G (2002) The simple fool's guide to PCR. Department of Zoology and Kewalo Marine Laboratory, University of Hawaii, Honolulu, Hawaii, 45 pp. https://stacks.stanford.edu/file/druid:yh-393jm6703/Simple_Fool%27s_Master%20PCR.pdf [accessed on 14 June 2021]
- Pante E, Simon-Bouhet B (2013) marmap: A package for importing, plotting and analyzing bathymetric and topographic data in R. *PLoS ONE* 8: e73051. <https://doi.org/10.1371/journal.pone.0073051>
- Pasulka AL, Goffredi SK, Tavormina PL, Dawson KS, Levin LA, Rouse GW, Orphan VJ (2017) Colonial tube-dwelling ciliates influence methane cycling and microbial diversity within methane seep ecosystems. *Frontiers in Marine Science* 3: 276. <https://doi.org/10.3389/fmars.2016.00276>
- Paterson GLJ, Baker AN (1988) A revision of the genus *Ophiambix* (Echinodermata: Ophiuroidea) including the description of a new species. *Journal of Natural History* 22: 1579–1590. <https://doi.org/10.1080/00222938800770991>
- Paull CK, Hecker B, Commeau R, Freeman-Lynde RP, Neumann C, Corso WP, Golubic S, Hook JE, Sikes E, Curray J (1984) Biological communities at the Florida Escarpment resemble hydrothermal vent taxa. *Science* 226: 965–967. <https://doi.org/10.1126/science.226.4677.965>
- Pawson DL (1965) The bathyal holothurians of the New Zealand region. *Zoology Publications from Victoria University of Wellington* 39: 1–33. <http://nzetc.victoria.ac.nz/tm/scholarly/tei-Vic39Zool-front.html> [accessed on 22 May 2020]
- Pawson DL (1970) 52 The marine fauna of New Zealand: Sea cucumbers (Echinodermata: Holothuroidea). New Zealand Department of Scientific and Industrial Research Bulletin 201. New Zealand Oceanographic Institute Memoir No. 52. New Zealand Oceanographic Institute, Wellington. <https://archive.org/details/nzoimemoir00521970> [accessed on 22 May 2020]
- Payne CY, Tilic E, Boschen-Rose RE, Gannon A, Stiller J, Hiley AS, Grupe BM, Mah CL, Rouse GW (2023) *Xyloplax princealberti* (Asteroidea, Echinodermata): A new species that is not always associated with wood falls. *Diversity* 15: 1212. <https://doi.org/10.3390/d15121212>
- Pearson KAM, Rouse GW (2022) Vampire Worms; A revision of *Galapagomystides* (Phyllodocidae, Annelida), with the description of three new species. *Zootaxa* 5128: 451–485. <https://doi.org/10.11646/zootaxa.5128.4.1>
- Peek AS, Gustafson RG, Lutz RA, Vrijenhoek RC (1997) Evolutionary relationships of deep-sea hydrothermal vent and cold-water seep clams (Bivalvia: Vesicomidae): Results from mitochondrial cytochrome oxidase subunit I. *Marine Biology* 130: 151–161. <https://doi.org/10.1007/s002270050234>

- Peek AS, Gaut BS, Feldman RA, Barry JP, Kochevar RE, Lutz RA, Vrijenhoek RC (2000) Neutral and nonneutral mitochondrial genetic variation in deep-sea clams from the family Vesicomidae. *Journal of Molecular Evolution* 50: 141–153. <https://doi.org/10.1007/s002399910016>
- Pereira OS, Gonzalez J, Mendoza GF, Le J, Coscino CL, Lee RW, Cortés J, Cordes EE, Levin LA (2021) The dynamic influence of methane seepage on macrofauna inhabiting authigenic carbonates. *Ecosphere* 12: e03744. <https://doi.org/10.1002/ecs2.3744>
- Pereira OS, Gonzalez J, Mendoza G, Le J, McNeill M, Ontiveros J, Lee RW, Rouse GW, Cortés J, Levin LA (2022) Does substrate matter in the deep sea? A comparison of bone, wood, and carbonate rock colonizers. *PLoS ONE* 17: e0271635. <https://doi.org/10.1371/journal.pone.0271635>
- Pereira OS, Jacobsen M, Carson R, Cortés J, Levin LA (2024) Understanding and valuing human connections to deep-sea methane seeps off Costa Rica. *Ecological Economics* 223: 108228. <https://doi.org/10.1016/j.ecolecon.2024.108228>
- Pérez-Barros P, Albano M, Diez MJ, Lovrich GA (2020) Pole to pole: the deep-sea king crab *Lithodes couesi* (Decapoda: Lithodidae) in the Burdwood Bank, Southwestern Atlantic Ocean. *Polar Biology* 43: 81–86. <https://doi.org/10.1007/s00300-019-02609-x>
- Petit RE (1983) A new species of *Cancellaria* (Mollusca: Cancellariidae) from the northern Gulf of Mexico. *Proceedings of the Biological Society of Washington* 96: 250–252. <https://www.biodiversitylibrary.org/item/107577> [accessed on 5 March 2021]
- Pettibone MH (1991) Polynoids commensal with gorgonian and stylasterid corals, with a new genus, new combinations, and new species (Polychaeta: Polynoidae: Polynoinae). *Proceedings of the Biological Society of Washington* 104: 688–713. <https://www.biodiversitylibrary.org/item/108199> [accessed on 12 April 2021]
- Pleijel F, Jondelius U, Norlinder E, Nygren A, Oxelman B, Schander C, Sundberg P, Thollesson M (2008) Phylogenies without roots? A plea for the use of vouchers in molecular phylogenetic studies. *Molecular Phylogenetics and Evolution* 48: 369–371. <https://doi.org/10.1016/j.ympev.2008.03.024>
- Portail M, Olu K, Escobar-Briones E, Caprais JC, Menot L, Waeles M, Cruaud P, Sarradin PM, Godfroy A, Sarrazin J (2015) Comparative study of vent and seep macrofaunal communities in the Guaymas Basin. *Biogeosciences Discussions* 12: 8497–8571. <https://doi.org/10.5194/bgd-12-8497-2015>
- Quattrini AM, Rodríguez E, Faircloth BC, Cowman PF, Brugler MR, Farfan GA, Hellberg ME, Kitahara MV, Morrison CL, Paz-García DA, Reimer JD, McFadden CS (2020) Palaeoclimate ocean conditions shaped the evolution of corals and their skeletons through deep time. *Nature Ecology and Evolution* 4: 1531–1538. <https://doi.org/10.1038/s41559-020-01291-1>
- Ramírez-Llodra E (2024) ChEssBase: an online information system on biodiversity and biogeography of deep-sea fauna from chemosynthetic ecosystems. ChEssBase. Version 2. <http://ipt.vliz.be/eurobis/resource?r=chessbase> [accessed on 4 April 2024]
- Ramírez Llodra E, Tyler PA, German CR (2003) Biogeography of deep-water chemosynthetic ecosystems (ChEss): Exploring the southern oceans. *Gayana* 67: 168–176. <https://doi.org/10.4067/S0717-65382003000200006>
- Ramírez-Llodra E, Tyler PA, Baker MC, Bergstad OA, Clark MR, Escobar E, Levin LA, Menot L, Rowden AA, Smith CR, van Dover CL (2011) Man and the last great wilderness: Human impact on the deep sea. *PLoS ONE* 6: e22588. <https://doi.org/10.1371/journal.pone.0022588>

- Regier JC, Shultz JW, Zwick A, Hussey A, Ball B, Wetzer R, Martin JW, Cunningham CW (2010) Arthropod relationships revealed by phylogenomic analysis of nuclear protein-coding sequences. *Nature* 463: 1079–1083. <https://doi.org/10.1038/nature08742>
- Reich A, Dunn C, Akasaka K, Wessel G (2015) Phylogenomic analyses of Echinodermata support the sister groups of Asterozoa and Echinozoa. *PLoS ONE* 10: e0119627. <https://doi.org/10.1371/journal.pone.0119627>
- Retamal MA, Guzmán G, De los Ríos-Escalante P (2020) An annotated checklist and bibliography of deep-water isopods and decapod crustaceans from Chile, including the submarine ridges Salas y Gomez and Nazca Plates. In: Hendrickx ME (Ed.), *Deep-Sea Pycnogonids and Crustaceans of the Americas*. Springer, Cham, 585–602. https://doi.org/10.1007/978-3-030-58410-8_22
- Reuscher M, Fiege D, Wehe T (2012) Terebellomorph polychaetes from hydrothermal vents and cold seeps with the description of two new species of Terebellidae (Annelida: Polychaeta) representing the first records of the family from deep-sea vents. *Journal of the Marine Biological Association of the United Kingdom* 92: 997–1012. <https://doi.org/10.1017/S0025315411000658>
- Ritger S, Carson B, Suess E (1987) Methane-derived authigenic carbonates formed by subduction-induced pore-water expulsion along the Oregon/Washington margin. *Geological Society of America Bulletin* 98: 147–156. [https://doi.org/10.1130/0016-7606\(1987\)98<147:MACFBS>2.0.CO;2](https://doi.org/10.1130/0016-7606(1987)98<147:MACFBS>2.0.CO;2)
- Robertson DR, Angulo A, Baldwin CC, Pitassy D, Driskell A, Weigt L, Navarro IJF (2017) Deep-water bony fishes collected by the B/O Miguel Oliver on the shelf edge of Pacific Central America: An annotated, illustrated and DNA-barcoded checklist. *Zootaxa* 4348: 1–125. <https://doi.org/10.11646/zootaxa.4348.1.1>
- Rodríguez-Flores PC, Seid CA, Rouse GW, Giribet G (2023) Cosmopolitan abyssal lineages? A systematic study of East Pacific deep-sea squat lobsters (Decapoda: Galatheoidea: Munidopsidae). *Invertebrate Systematics* 37: 14. <https://doi.org/10.1071/IS22030>
- Rojas-Jimenez K, Grossart HP, Cordes E, Cortés J (2020) Fungal communities in sediments along a depth gradient in the Eastern Tropical Pacific. *Frontiers in Microbiology* 11: 575207. <https://doi.org/10.3389/fmicb.2020.575207>
- Rouse GW, Kupriyanova EK (2021) *Laminatubus* (Serpulidae, Annelida) from eastern Pacific hydrothermal vents and methane seeps, with description of two new species. *Zootaxa* 4915: 1–27. <https://doi.org/10.11646/zootaxa.4915.1.1>
- Rouse GW, Goffredi SK, Vrijenhoek RC (2004) *Osedax*: Bone-eating marine worms with dwarf males. *Science* 305: 668–671. <https://doi.org/10.1126/science.1098650>
- Rouse GW, Carvajal JI, Pleijel F (2018) Phylogeny of Hesionidae (Aciculata, Annelida), with four new species from deep-sea eastern Pacific methane seeps, and resolution of the affinity of *Hesiolyra*. *Invertebrate Systematics* 32: 1050–1068. <https://doi.org/10.1071/IS17092>
- Rouse GW, Pleijel F, Tilic E (2022) *Annelida*. Oxford University Press, United Kingdom, 432 pp. <https://doi.org/10.1093/oso/9780199692309.001.0001>
- Roux M (2004) New hyocrinid crinoids (Echinodermata) from submersible investigations in the Pacific Ocean. *Pacific Science* 58: 597–613. <https://doi.org/10.1353/psc.2004.0042>
- Rybakova E, Krylova E, Mordukhovich V, Galkin S, Alalykina I, Smirnov I, Sanamyan N, Nekhaev I, Vinogradov G, Shilov V, Prudkovsky A, Kolpakov E, Gebruk A, Adrianov A (2022) Methane seep communities on the Koryak slope in the Bering Sea. *Deep Sea Research Part II: Topical Studies in Oceanography* 206: 105203. <https://doi.org/10.1016/j.dsr2.2022.105203>

- Rybakova E, Krylova E, Mordukhovich V, Galkin S, Alalykina I, Sanamyan N, Nekhaev I, Vinogradov G, Shilov V, Pakhnevich A, Gebruk A, Adrianov A (2023) Mega- and macrofauna of the hydrothermally active submarine Piip Volcano (the southwestern Bering Sea). *Deep Sea Research Part II: Topical Studies in Oceanography* 208: 105268. <https://doi.org/10.1016/j.dsr2.2023.105268>
- Saeedi H, Bernardino AF, Shimabukuro M, Falchetto G, Sumida PYG (2019) Macrofaunal community structure and biodiversity patterns based on a wood-fall experiment in the deep South-west Atlantic. *Deep Sea Research Part I: Oceanographic Research Papers* 145: 73–82. <https://doi.org/10.1016/j.dsr.2019.01.008>
- Sagorny C, von Döhren J, Rouse GW, Tilic E (2022) Cutting the ribbon: bathyal Nemertea from seeps along the Costa Rica margin, with descriptions of 2 new genera and 9 new species. *European Journal of Taxonomy* 845: 132–174. <https://doi.org/10.5852/ejt.2022.845.1959>
- Sahling H, Masson DG, Ranero CR, Hühnerbach V, Weinrebe W, Klaucke I, Bürk D, Brückmann W, Suess E (2008) Fluid seepage at the continental margin offshore Costa Rica and southern Nicaragua. *Geochemistry, Geophysics, Geosystems* 9: Q05S05. <https://doi.org/10.1029/2008GC001978>
- Salazar-Vallego SI (2012) Revision of *Flabelligera* Sars, 1829 (Polychaeta: Flabelligeridae). *Zootaxa* 3203: 1–64. <https://doi.org/10.11646/zootaxa.3203.1.1>
- Salazar-Vallejo SI (2017) Revision of *Brada* Stimpson, 1853, and *Bradabyssa* Hartman, 1967 (Annelida, Flabelligeridae). *Zootaxa* 4343: 1–98. <https://doi.org/10.11646/zootaxa.4343.1.1>
- Salazar-Vallejo SI (2020a) Four new deep-water flabelligerid species from Pacific Costa Rica (Annelida, Sedentaria, Flabelligeridae). *Zootaxa* 4885: 560–578. <https://doi.org/10.11646/zootaxa.4885.4.6>
- Salazar-Vallejo SI (2020b) Revision of *Leocrates* Kinberg, 1866 and *Leocratides* Ehlers, 1908 (Annelida, Errantia, Hesionidae). *Zootaxa* 4739: 1–114. <https://doi.org/10.11646/zootaxa.4739.1.1>
- Salinas-de-León P, Martí-Puig P, Buglass S, Arnés-Urgellés C, Rastoin-Laplane E, Creemers M, Cairns S, Fisher C, O'Hara T, Ott B, Raineault NA, Reiswig H, Rouse G, Rowley S, Shank TM, Suarez J, Watling L, Wicksten MK, Marsh L (2020) Characterization of deep-sea benthic invertebrate megafauna of the Galapagos Islands. *Scientific Reports* 10: 1–13. <https://doi.org/10.1038/s41598-020-70744-1>
- Santagata S, Tunnell Jr JW (2009) Brachiopoda of the Gulf of Mexico. In: Felder DL, Camp DK (Eds), *Gulf of Mexico—Origin, Waters, and Biota*. Texas A&M University Press, College Station, TX, 1137–1141. <https://muse.jhu.edu/book/344> [accessed on 23 March 2023]
- Sasaki T, Warén A, Kano Y, Okutani T, Fujikura K (2010) Gastropods from recent hot vents and cold seeps: systematics, diversity and life strategies. In: Kiel S (Ed.), *The Vent and Seep Biota*. Topics in Geobiology. Springer, Dordrecht, 169–254. https://doi.org/10.1007/978-90-481-9572-5_7
- Schwabe E, Sellanes J (2004) A new species of *Lepidozona* (Mollusca: Polyplacophora: Ischnochitonidae), found on whale bones off the coast of Chile. *Iberus* 22: 147–153. <https://doi.org/10.5281/zenodo.4516397>
- Schwabe E, Sellanes J (2010) Revision of Chilean bathyal chitons (Mollusca: Polyplacophora) associated with cold-seeps, including description of a new species of *Leptochiton* (Leptochitonidae). *Organisms Diversity & Evolution* 10: 31–55. <https://doi.org/10.1007/s13127-009-0002-6>
- Schwentner M, Richter S, Rogers DC, Giribet G (2018) Tetraconatan phylogeny with special focus on Malacostraca and Branchiopoda: Highlighting the strength of

- taxon-specific matrices in phylogenomics. *Proceedings of the Royal Society B: Biological Sciences* 285: 20181524. <https://doi.org/10.1098/rspb.2018.1524>
- Selig GM, Netburn AN, Malik M (2019) Distributions of the pelagic holothurian *Pelagothuria* in the Central Pacific Ocean as observed by remotely-operated vehicle surveys. *Frontiers in Marine Science* 6: 684. <https://doi.org/10.3389/fmars.2019.00684>
- Sellanes J, Quiroga E, Neira C (2008) Megafauna community structure and trophic relationships at the recently discovered Concepción Methane Seep Area, Chile, ~36°S. *ICES Journal of Marine Science* 65: 1102–1111. <https://doi.org/10.1093/icesjms/fsn099>
- Shen Z, Koch NM, Seid CA, Tilic E, Rouse GW (2024) Three new species of deep-sea wood-associated sea stars (Asteroidea: Caymanostellidae) from the Eastern Pacific. *Zootaxa* 5536: 351–388. <https://doi.org/10.11646/zootaxa.5536.3.1>
- Sibuet M, Olu K (1998) Biogeography, biodiversity and fluid dependence of deep-sea cold-seep communities at active and passive margins. *Deep Sea Research Part II: Topical Studies in Oceanography* 45: 517–567. [https://doi.org/10.1016/S0967-0645\(97\)00074-X](https://doi.org/10.1016/S0967-0645(97)00074-X)
- Sigovini M, Keppel E, Tagliapietra D (2016) Open Nomenclature in the biodiversity era. *Methods in Ecology and Evolution* 7: 1217–1225. <https://doi.org/10.1111/2041-210X.12594>
- Sigwart JD, Chen C, Tilic E, Vences M, Riehl T (2023) Why is there no service to support taxonomy? *BioEssays* 45: e2300070. <https://doi.org/10.1002/bies.202300070>
- Simoneit BRT, Lonsdale PF, Edmond JM, Shanks WC (1990) Deep-water hydrocarbon seeps in Guaymas Basin, Gulf of California. *Applied Geochemistry* 5: 41–49. [https://doi.org/10.1016/0883-2927\(90\)90034-3](https://doi.org/10.1016/0883-2927(90)90034-3)
- Sirenko B, Sellanes J (2016) Update of the genus *Leptochiton* (Mollusca: Polyplacophora) in Chilean deep waters: Three new reports and description of two new species. *Zootaxa* 4173: 259–279. <https://doi.org/10.11646/zootaxa.4173.3.5>
- Sirenko B, Saito H, Schwabe E (2022) A redescription of *Leptochiton belknapi* Dall, 1878 (Mollusca: Polyplacophora: Leptochitonidae), the type species of the new genus *Belknapchiton*. *Zootaxa* 5205: 101–124. <https://doi.org/10.11646/zootaxa.5205.2.1>
- Sjölin E, Erséus C, Källersjö M (2005) Phylogeny of Tubificidae (Annelida, Clitellata) based on mitochondrial and nuclear sequence data. *Molecular Phylogenetics and Evolution* 35: 431–441. <https://doi.org/10.1016/j.ympev.2004.12.018>
- Smith SI (1885) On some new or little known decapod Crustacea, from recent Fish Commission dredgings off the east coast of the United States. *Proceedings of the United States National Museum* 7: 493–511. <https://doi.org/10.5479/si.00963801.455.493>
- Smith CR, Baco AR (2003) Ecology of whale falls at the deep-sea floor. In: Gibson RN, Atkinson RJA (Eds), *Oceanography and Marine Biology: An Annual Review*. Taylor & Francis, 311–354. <https://www.routledge.com/Oceanography-and-Marine-Biology-An-Annual-Review-Volume-41-An-Annual/Gibson-Atkinson/p/book/9780415254632> [accessed on 25 February 2021]
- Smith CR, Kukert H, Wheatcroft RA, Jumars PA, Deming JW (1989) Vent fauna on whale remains. *Nature* 341: 27–28. <https://doi.org/10.1038/341027a0>
- Smith SA, Wilson NG, Goetz FE, Feehery C, Andrade SCS, Rouse GW, Giribet G, Dunn CW (2011) Resolving the evolutionary relationships of molluscs with phylogenomic tools. *Nature* 480: 364–367. <https://doi.org/10.1038/nature10526>
- Solís-Weiss V (1993) *Grassleia hydrothermalis*, a new genus and species of Ampharetidae (Annelida: Polychaeta) from the hydrothermal vents off the Oregon coast (U.S.A.), at Gorda Ridge. *Proceedings of the Biological Society of Washington* 106: 661–665. <https://www.biodiversitylibrary.org/part/44993> [accessed on 15 April 2021]

- Solís-Weiss V, Hernández-Alcántara P (1994) *Amphisamytha fauchaldi*: A new species of ampharetid (Annelida: Polychaeta) from the hydrothermal vents at Guaymas Basin, Mexico. *Bulletin of the Southern California Academy of Sciences* 93: 127–134. <https://www.biodiversitylibrary.org/part/42351> [accessed on 5 May 2020]
- Solís-Weiss V, Hilbig B (1992) Redescription of *Ophryotrocha platycephale* Blake (Polychaeta, Dorvilleidae) from the Guaymas Basin hydrothermal vents. *Bulletin of Southern California Academy of Sciences* 92: 92–96. <https://www.biodiversitylibrary.org/item/106555> [accessed on 14 July 2022]
- Song X, Lyu M, Zhang X, Ruthensteiner B, Ahn I-Y, Pastorino G, Wang Y, Gu Y, Ta K, Sun J, Liu X, Han J, Ke C, Peng X (2021) Large plastic debris dumps: New biodiversity hot spots emerging on the deep-sea floor. *Environmental Science & Technology Letters* 8: 148–154. <https://doi.org/10.1021/acs.estlett.0c00967>
- Southward EC, Schulze A, Tunnicliffe V (2002) Vestimentiferans (Pogonophora) in the Pacific and Indian Oceans: A new genus from Lihir Island (Papua New Guinea) and the Java Trench, with the first report of *Arcovestia ivanovi* from the North Fiji Basin. *Journal of Natural History* 36: 1179–1197. <https://doi.org/10.1080/00222930110040402>
- Squires HJ, Barragan JH (1976) A new species of *Plesionika* (Crustacea, Decapoda, Pandalidae) from the Pacific coast of Colombia. *Pacific Science* 30: 113–117. <http://hdl.handle.net/10125/10771> [accessed on 22 August 2022]
- Stebbing TRR (1888) Report on the Amphipoda collected by H.M.S. Challenger during the years 1873–1876. *Zoology Volume XXIX. Part LXVII*. In: Report on the scientific results of the voyage of H.M.S. Challenger during the years 1873–76 under the command of Captain George S. Nares, RN, FRS, and the late Captain Frank Tourle Thomson, R.N. Neill, Edinburgh, 1–1737. <https://doi.org/10.5962/bhl.title.6513>
- Stebbins TD, Eernisse DJ (2009) Chitons (Mollusca: Polyplacophora) known from benthic monitoring programs in the Southern California Bight. *The Festivus* 41: 53–100. <https://www.biodiversitylibrary.org/page/49523140> [accessed on 3 June 2020]
- Stepanov VG, Panina EG (2016) A checklist of the holothurians of the far eastern seas of Russia. *SPC Beche-de-mer Information Bulletin* 36: 24–35. <https://purl.org/spc/digilib/doc/cztyw> [accessed on 27 May 2021]
- Stiller J, Rousset V, Pleijel F, Chevalloné P, Vrijenhoek RC, Rouse GW (2013) Phylogeny, biogeography and systematics of hydrothermal vent and methane seep *Amphisamytha* (Ampharetidae, Annelida), with descriptions of three new species. *Systematics and Biodiversity* 11: 35–65. <https://doi.org/10.1080/14772000.2013.772925>
- Stiller J, Tilic E, Rousset V, Pleijel F, Rouse GW (2020) Spaghetti to a tree: A robust phylogeny for Terebelliformia (Annelida) based on transcriptomes, molecular and morphological data. *Biology* 9: 1–28. <https://doi.org/10.3390/biology9040073>
- Stöhr S, Sautya S, Ingole B (2012) Brittle stars (Echinodermata: Ophiuroidea) from seamounts in the Andaman Sea (Indian Ocean): First account, with descriptions of new species. *Journal of the Marine Biological Association of the United Kingdom* 92: 1195–1208. <https://doi.org/10.1017/S0025315412000240>
- Strand M, Norenburg J, Alfaya JE, Ángel Fernández-Álvarez F, Andersson HS, Andrade SCS, Bartolomaeus T, Beckers P, Bigatti G, Cherneva I, Chernyshev A, Chung BM, Döhren J, Giribet G, Gonzalez-Cueto J, Herrera-Bachiller A, Hiebert T, Hookabe N, Junoy J, Kajihara H, Krämer D, Kvist S, Magarlamov TY, Maslakova S, Mendes CB, Okazaki R, Sagorny C, Schwartz M, Sun S, Sundberg P, Turbeville JM, Xu C (2018) Nemeritean taxonomy—Implementing changes in the higher ranks, dismissing Anopla and Enopla. *Zoologica Scripta* 48: 118–119. <https://doi.org/10.1111/zsc.12317>

- Suarez AV, Tsutsui ND (2004) The value of museum collections for research and society. *Bioscience* 54: 66–74. [https://doi.org/10.1641/0006-3568\(2004\)054\[0066:T-VOMCF\]2.0.CO;2](https://doi.org/10.1641/0006-3568(2004)054[0066:T-VOMCF]2.0.CO;2)
- Suárez-Mozo NY, Hendrickx ME (2016) New record for the deep-sea genus *Tripoplax* (Mollusca: Polyplacophora) in the eastern Pacific. *Marine Biodiversity Records* 9: 10. <https://doi.org/10.1186/s41200-016-0011-z>
- Suárez-Mozo NY, Valentich-Scott P, Hendrickx ME (2019) Deep-water bivalves from the oxygen minimum zone area off the western Peninsula of Baja California, Mexico. *Molluscan Research* 39: 99–109. <https://doi.org/10.1080/13235818.2018.1508550>
- Suess E, Carson B, Ritger S, Moore J, Jones M, Kulm L, Cochrane G (1985) Biological communities at vent sites along the subduction zone off Oregon. *Bulletin of the Biological Society of Washington* 6: 475–484. <https://oceanrep.geomar.de/id/eprint/31574> [accessed 12 July 2021]
- Summers MM, Messing CG, Rouse GW (2014a) Phylogeny of Comatulidae (Echinodermata: Crinoidea: Comatulida): A new classification and an assessment of morphological characters for crinoid taxonomy. *Molecular Phylogenetics and Evolution* 80: 319–339. <https://doi.org/10.1016/j.ympev.2014.06.030>
- Summers MM, Al-Hakim II, Rouse GW (2014b) Turbo-taxonomy: 21 new species of Myzostomida (Annelida). *Zootaxa* 3873: 301–344. <https://doi.org/10.11646/zootaxa.3873.4.1>
- Summers M, Pleijel F, Rouse GW (2015) Whale falls, multiple colonisations of the deep, and the phylogeny of Hesionidae (Annelida). *Invertebrate Systematics* 29: 105–123. <https://doi.org/10.1071/IS14055>
- Tapia-Guerra JM, Mecho A, Easton EE, Gallardo M de LÁ, Gorny M, Sellanes J (2021) First description of deep benthic habitats and communities of oceanic islands and seamounts of the Nazca Desventuradas Marine Park, Chile. *Scientific Reports* 11: 6209. <https://doi.org/10.1038/s41598-021-85516-8>
- Tavormina PL, Ussler W, Steele JA, Connon SA, Klotz MG, Orphan VJ (2013) Abundance and distribution of diverse membrane-bound monooxygenase (Cu-MMO) genes within the Costa Rica oxygen minimum zone. *Environmental Microbiology Reports* 5: 414–423. <https://doi.org/10.1111/1758-2229.12025>
- Templado J, Paulay G, Gittenberger A, Meyer C (2010) Chapter 11 Sampling the Marine Realm. In: Eymann J, Degreaf J, Häuser C, Monje JC, Samyn Y, Vandenspiegel D (Eds), *Manual on Field Recording Techniques and Protocols for All Taxa Biodiversity Inventories*. *Abc Taxa* 8: 273–307. <https://abctaxa.naturalsciences.be/pdf-8/> [accessed on 16 July 2019]
- Thomas EA, Liu R, Amon D, Copley JT, Glover AG, Helyar SJ, Olu K, Wiklund H, Zhang H, Sigwart JD (2020) *Chiridota heheva*—the cosmopolitan holothurian. *Marine Biodiversity* 50: 110. <https://doi.org/10.1007/s12526-020-01128-x>
- Thomson CW (1873) *The Depths of the Sea. An Account of the General Results of the Dredging Cruises of the H.M.S.S. "Porcupine" and "Lightning" During the Summers of 1868, 1869, and 1870, Under the Scientific Direction of Dr. Carpenter, F.R.S., J. Gwyn Jeffreys, F.R.S. and Dr. Wyville Thomson, F.R.S.* Macmillan and Co., London, 527 pp. <https://doi.org/10.5962/bhl.title.19314>
- Thurber AR, Jones WJ, Schnabel K (2011) Dancing for food in the deep sea: Bacterial farming by a new species of Yeti crab. *PLoS ONE* 6: 1–12. <https://doi.org/10.1371/journal.pone.0026243>
- Thurber AR, Levin LA, Orphan VJ, Marlow JJ (2012) Archaea in metazoan diets: Implications for food webs and biogeochemical cycling. *The ISME Journal* 6: 1602–1612. <https://doi.org/10.1038/ismej.2012.16>

- Tunnicliffe V, McArthur AG, McHugh D (1998) A biogeographical perspective of the deep-sea hydrothermal vent fauna. *Advances in Marine Biology* 34: 353–442. [https://doi.org/10.1016/S0065-2881\(08\)60213-8](https://doi.org/10.1016/S0065-2881(08)60213-8)
- Tunnicliffe V, Juniper SK, Sibuet M (2003) Reducing environments of the deep-sea floor. In: Tyler PA (Ed.) *Ecosystems of the Deep Oceans*. Elsevier Press, Amsterdam, 81–110.
- Tunnicliffe V, Chen C, Giguère T, Rowden AA, Watanabe HK, Brunner O (2024) Hydrothermal vent fauna of the western Pacific Ocean: Distribution patterns and biogeographic networks. *Diversity and Distributions* 30: e13794. <https://doi.org/10.1111/ddi.13794>
- Turnipseed M, Knick KE, Lipcius RN, Dreyer J, Van Dover CL (2003) Diversity in mussel beds at deep-sea hydrothermal vents and cold seeps. *Ecology Letters* 6: 518–523. <https://doi.org/10.1046/j.1461-0248.2003.00465.x>
- Uribe JE, Zardoya R, Puillandre N (2018) Phylogenetic relationships of the conoidean snails (Gastropoda: Caenogastropoda) based on mitochondrial genomes. *Molecular Phylogenetics and Evolution* 127: 898–906. <https://doi.org/10.1016/j.ympev.2018.06.037>
- Uribe JE, González VL, Irisarri I, Kano Y, Herbert DG, Strong EE, Harasewych MG (2022) A phylogenomic backbone for gastropod molluscs. *Systematic Biology* 71: 1271–1280. <https://doi.org/10.1093/sysbio/syac045>
- Valentich-Scott P (1998) Class Bivalvia. In: Valentich-Scott P, Blake JA (Eds), Volume 8: The Mollusca Part 1, The Aplacophora, Polyplacophora, Scaphopoda, Bivalvia, and Cephalopoda. *Taxonomic Atlas of the Benthic Fauna of the Santa Maria Basin and the Western Santa Barbara Channel*. Santa Barbara Museum of Natural History, CA, USA, 97–173.
- Valentich-Scott P, Coan EV, Zelaya DG (2020) *Bivalve Seashells of Western South America*. 1st edn. Santa Barbara Museum of Natural History, CA, USA.
- Van Dover CL, German CR, Speer KG, Parson LM, Vrijenhoek RC (2002) Evolution and biogeography of deep-sea vent and seep invertebrates. *Science* 295: 1253–1257. <https://doi.org/10.1126/science.1067361>
- Van Dover CL, Smith CR, Ardrón J, Dunn D, Gjerde K, Levin L, Smith S, Arnaud-Haond S, Beaudoin Y, Bezaury J, Boland G, Billett D, Carr M, Cherkashov G, Cook A, DeLeo F, Fisher CR, Godet L, Halpin P, Lodge M, Menot L, Miller K, Naudts L, Nugent C, Pendleton L, Plouviez S, Rowden AA, Santos RS, Shank T, Tao C, Tawake A, Thurnherr A, Treude T (2012) Designating networks of chemosynthetic ecosystem reserves in the deep sea. *Marine Policy* 36: 378–381. <https://doi.org/10.1016/j.marpol.2011.07.002>
- Vedenin AA, Kokarev VN, Chikina MV, Basin AB, Galkin SV, Gebruk AV (2020) Fauna associated with shallow-water methane seeps in the Laptev Sea. *PeerJ* 8: e9018. <https://doi.org/10.7717/peerj.9018>
- Villalobos-Guerrero TF, Huč S, Tilic E, Hiley AS, Rouse GW (2024) A remarkable new deep-sea nereidid (Annelida: Nereididae) with gills. *PLoS ONE* 19: e0297961. <https://doi.org/10.1371/journal.pone.0297961>
- von Huene R, Ranero CR, Weinrebe W, Hinz K (2000) Quaternary convergent margin tectonics of Costa Rica, segmentation of the Cocos Plate, and Central American volcanism. *Tectonics* 19: 314–334. <https://doi.org/10.1029/1999TC001143>
- Wagner D, van der Meer L, Gorny M, Sellanes J, Gaymer CF, Soto EH, Easton EE, Friedlander AM, Lindsay DJ, Molodtsova TN, Boteler B, Durussel C, Gjerde KM, Currie D, Gianni M, Brooks CM, Shiple MJ, Wilhelm TA, Quesada M, Thomas T, Dunstan PK, Clark NA, Villanueva LA, Pyle RL, Clark MR, Georgian SE, Morgan LE (2021) The Salas y Gómez and Nazca ridges: A review of the importance, opportunities and challenges for protecting a global diversity hotspot on the high seas. *Marine Policy* 126: 104377. <https://doi.org/10.1016/j.marpol.2020.104377>

- Warén A, Bouchet P (1986) Four new species of *Provanna* Dall (Prosobranchia, Cerithiacea?) from East Pacific hydrothermal sites. *Zoologica Scripta* 15: 157–164. <https://doi.org/10.1111/j.1463-6409.1986.tb00218.x>
- Warén A, Bouchet P (1989) New gastropods from East Pacific hydrothermal vents. *Zoologica Scripta* 18: 67–102. <https://doi.org/10.1111/j.1463-6409.1989.tb00124.x>
- Warén A, Bouchet P (1993) New records, species, genera, and a new family of gastropods from hydrothermal vents and hydrocarbon seeps. *Zoologica Scripta* 22: 1–90. <https://doi.org/10.1111/j.1463-6409.1993.tb00342.x>
- Warén A, Bouchet P (2001) Gastropoda and Monoplacophora from hydrothermal vents and seeps; new taxa and records. *The Veliger* 44: 116–231. <https://www.biodiversitylibrary.org/part/97997> [accessed on 11 June 2020]
- Warén A, Bouchet P (2009) New gastropods from deep-sea hydrocarbon seeps off West Africa. *Deep Sea Research Part II: Topical Studies in Oceanography* 56: 2326–2349. <https://doi.org/10.1016/j.dsr2.2009.04.013>
- Warén A, Ponder WF (1991) New species, anatomy, and systematic position of the hydrothermal vent and hydrocarbon seep gastropod family Provannidae fam. n. (Caenogastropoda). *Zoologica Scripta* 20: 27–56. <https://doi.org/10.1111/j.1463-6409.1991.tb00273.x>
- Warén A, Rouse GW (2016) A new genus and species of Cataegidae (Gastropoda: Seguenzioidea) from eastern Pacific Ocean methane seeps. *Novapex* 17: 59–66. <https://www.biodiversitylibrary.org/part/283777> [accessed on 1 February 2017]
- Watanabe H, Fujikura K, Kojima S, Miyazaki J-I, Fujiwara Y (2010) Japan: Vents and seeps in close proximity. In: Kiel S (Ed.), *The Vent and Seep Biota*. Springer, Dordrecht, 379–401. https://doi.org/10.1007/978-90-481-9572-5_12
- Watson C, Ignacio Carvajal J, Sergeeva NG, Pleijel F, Rouse GW (2016) Free-living calamyzin chrysopetalids (Annelida) from methane seeps, anoxic basins, and whale falls. *Zoological Journal of the Linnean Society* 177: 700–719. <https://doi.org/10.1111/zoj.12390>
- Webb M (1969) *Lamellibrachia barhami*, gen. nov., sp. nov. (Pogonophora), from the northeast Pacific. *Bulletin of Marine Science* 19: 18–47. <http://www.ingentaconnect.com/content/umrsmas/bullmar/1969/00000019/00000001/art00002> [accessed on 4 May 2020]
- Wehrtmann IS, Cortés J (Eds) (2009a) *Marine Biodiversity of Costa Rica, Central America*. Springer, Dordrecht. <https://doi.org/10.1007/978-1-4020-8278-8>
- Wehrtmann IS, Cortés J (2009b) Contents of CD-ROM. In: Wehrtmann IS, Cortés J (Eds), *Marine Biodiversity of Costa Rica, Central America*. Monographiae Biologicae. Springer, Dordrecht, 500 pp. <https://extras.springer.com/?query=978-1-4020-8277-1> [accessed on 22 May 2020]
- Wehrtmann IS, Echeverría-Sáenz S (2007) Crustacean fauna (Stomatopoda: Decapoda) associated with the deepwater fishery of *Heterocarpus vicarius* (Decapoda: Pandalidae) along the Pacific coast of Costa Rica. *Revista de Biología Tropical* 55 (Supplement 1): 121–130. <https://doi.org/10.15517/rbt.v55i0.5812>
- Wehrtmann IS, Herrera-Correal J, Vargas R, Hernández P (2010) Squat lobsters (Decapoda: Anomura: Galatheididae) from deepwater Pacific Costa Rica: species diversity, spatial and bathymetric distribution. *Nauplius* 18: 69–77. http://crustacea.org.br/wp-content/uploads/2014/02/nauplius-v18n1a07.Wehrtmann.et_al.pdf [accessed on 13 April 2020]
- Weis A, Melzer RR (2012) How did sea spiders recolonize the Chilean fjords after glaciation? DNA barcoding of Pycnogonida, with remarks on phylogeography of *Achelia assimilis* (Haswell, 1885). *Systematics and Biodiversity* 10: 361–374. <https://doi.org/10.1080/14772000.2012.716462>

- Wicksten MK (1980) New records of *Stereomastis sculpta pacifica* (Faxon) (Decapoda: Polychelidae) in the eastern Pacific Ocean. *Proceedings of the Biological Society of Washington* 93: 914–919. <https://www.biodiversitylibrary.org/part/45616> [accessed on 7 May 2020]
- Wicksten MK (1989) Ranges of offshore decapod crustaceans in the eastern Pacific Ocean. *Transactions of the San Diego Society of Natural History* 21: 291–316. <https://doi.org/10.5962/bhl.part.24590>
- Wicksten MK (2020) Lower slope and abyssal benthic decapods of the eastern Pacific. In: Hendrickx ME (Ed.) *Deep-Sea Pycnogonids and Crustaceans of the Americas*. Springer, Cham, 395–420. https://doi.org/10.1007/978-3-030-58410-8_17
- Wicksten MK, Méndez M (1982) New records and new species of the genus *Lebbeus* (Caridea: Hippolytidae) in the eastern Pacific Ocean. *Bulletin of Southern California Academy of Sciences* 81: 106–120. <https://www.biodiversitylibrary.org/part/43130> [accessed on 3 May 2021]
- Wiklund H, Altamira IV, Glover AG, Smith CR, Baco AR, Dahlgren TG (2012) Systematics and biodiversity of *Ophryotrocha* (Annelida, Dorvilleidae) with descriptions of six new species from deep-sea whale-fall and wood-fall habitats in the north-east Pacific. *Systematics and Biodiversity* 10: 243–259. <https://doi.org/10.1080/14772000.2012.693970>
- Williams AB (1988) New marine decapod crustaceans from waters influenced by hydrothermal discharge, brine, and hydrocarbon seepage. *Fishery Bulletin* 86: 263–287. <https://repository.si.edu/handle/10088/21612> [accessed on 24 May 2021]
- Williams GC (2011) The global diversity of sea pens (Cnidaria: Octocorallia: Pennatulacea). *PLoS ONE* 6: e22747. <https://doi.org/10.1371/journal.pone.0022747>
- Wilson EB (1881) Report on the Pycnogonida. Reports on the results of dredging, under the supervision of A. Agassiz, along the East Coast of the United States during the summer of 1880, by the United States Coast Survey steamer, “Blake,” Commander, J. R. Bartlett, U. S. N. *Bulletin of the Museum of Comparative Zoology at Harvard College* 8: 239–256. <https://www.biodiversitylibrary.org/page/28868670> [accessed on 19 May 2021]
- Winfield I, Hendrickx ME, Ortiz M (2020) Updated checklist of deep-sea amphipods (Amphilochidea and Senticaudata) from Western Mexico, NE Pacific Ocean. In: Hendrickx ME (Ed.), *Deep-Sea Pycnogonids and Crustaceans of the Americas*. Springer, Cham, 127–141. https://doi.org/10.1007/978-3-030-58410-8_5
- Wolfe JM, Breinholt JW, Crandall KA, Lemmon AR, Lemmon EM, Timm LE, Siddall ME, Bracken-Grissom HD (2019) A phylogenomic framework, evolutionary timeline and genomic resources for comparative studies of decapod crustaceans. *Proceedings of the Royal Society B: Biological Sciences* 286: 1–10. <https://doi.org/10.1098/rspb.2019.0079>
- WoRMS Editorial Board (2024) World Register of Marine Species. <https://doi.org/10.14284/170>
- Xu T, Sun J, Lv J, Kayama Watanabe H, Li T, Zou W, Rouse GW, Wang S, Qian PY, Bao Z, Qiu JW (2017) Genome-wide discovery of single nucleotide polymorphisms (SNPs) and single nucleotide variants (SNVs) in deep-sea mussels: Potential use in population genomics and cross-species application. *Deep Sea Research Part II: Topical Studies in Oceanography* 137: 318–326. <https://doi.org/10.1016/j.dsr2.2016.03.011>
- Yapur-Pancorvo AL, Quispe-Machaca M, Guzmán-Rivás F, Urzúa Á, Espinoza P (2023) The red squat lobster *Pleuroncodes monodon* in the Humboldt Current System: From their ecology to commercial attributes as marine bioresource. *Animals* 13: 2279. <https://doi.org/10.3390/ani13142279>
- Yen NK, Rouse GW (2020) Phylogeny, biogeography and systematics of Pacific vent, methane seep, and whale-fall *Parougia* (Dorvilleidae: Annelida), with eight new species. *Invertebrate Systematics* 34: 200–233. <https://doi.org/10.1071/IS19042>

- Zanol J, Carrera-Parra LF, Steiner TM, Amaral ACZ, Wiklund H, Ravara A, Budaeva N (2021) The current state of Eunicida (Annelida) systematics and biodiversity. *Diversity* 13: 1–54. <https://doi.org/10.3390/d13020074>
- Zapata-Hernández G, Sellanes J, Thurber AR, Levin LA (2014) Trophic structure of the bathyal benthos at an area with evidence of methane seep activity off southern Chile (~45°S). *Journal of the Marine Biological Association of the United Kingdom* 94: 659–669. <https://doi.org/10.1017/S0025315413001914>
- Zeidler W (2021) Review of the hyperiidean amphipod family Lycaeidae Claus, 1879 (Crustacea: Amphipoda: Hyperiidea). *Zootaxa* 5081: 1–59. <https://doi.org/10.11646/zootaxa.5081.1.1>
- Zhang Z, Schwartz S, Wagner L, Miller W (2000) A greedy algorithm for aligning DNA sequences. *Journal of Computational Biology* 7: 203–214. <https://doi.org/10.1089/10665270050081478>
- Zhang R, Fau M, Mah C, Eléaume M, Zhang D, Zhou Y, Lu B, Wang C (2024) Systematics of deep-sea starfish order Brisingida (Echinodermata: Asteroidea), with a revised classification and assessments of morphological characters. *Molecular Phylogenetics and Evolution* 191: 107993. <https://doi.org/10.1016/j.ympev.2023.107993>
- Zhong Z, Lan Y, Chen C, Zhou Y, Linse K, Li R, Sun J (2022) New mitogenomes in deep-water endemic Cocculinida and Neomphalida shed light on lineage-specific gene orders in major gastropod clades. *Frontiers in Ecology and Evolution* 10: 973485. <https://doi.org/10.3389/fevo.2022.973485>
- Zuleger E, Gieskes JM, You C-F (1996) Interstitial water chemistry of sediments of the Costa Rica accretionary complex off the Nicoya Peninsula. *Geophysical Research Letters* 23: 899–902. <https://doi.org/10.1029/96GL00386>

Supplementary material 1

List of all taxa occurring at the CRM seeps and new records in this work

Authors: Charlotte A. Seid, Avery S. Hiley, Marina F. McCowin, José I. Carvajal, Harim Cha, Shane T. Ah Yong, Oliver S. Ashford, Odalysa Breedy, Douglas J. Eernisse, Shana K. Goffredi, Michel E. Hendrickx, Kevin M. Kocot, Christopher L. Mah, Allison K. Miller, Nicolás Mongiardino Koch, Rich Mooi, Timothy D. O'Hara, Fredrik Pleijel, Josefín Stiller, Ekin Tilic, Paul Valentich-Scott, Anders Warén, Mary K. Wicksten, Nerida G. Wilson, Erik E. Cordes, Lisa A. Levin, Jorge Cortés, Greg W. Rouse

Data type: xlsx

Explanation note: For described species: depth range at the CRM, known biogeographic range, known occurrences at vents or organic falls, and new records are indicated. If only a single depth is reported, the depth value is shown as a minimum. Exact collection depths are provided where possible; approximations are indicated by ~. This table summarizes the taxonomic section of the main text, including newly published GenBank sequences, but introduces no new information.

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Link: <https://doi.org/10.3897/zookeys.1222.134385.suppl1>