

Increasing Information Content and Diagnosability in Family-Level Classifications

MATJAŽ KUNTNER^{1,2,3,4,5,*}, KLEMEN ČANDEK^{1,4}, MATJAŽ GREGORIČ², EVA TURK^{4,6}, CHRIS A. HAMILTON⁷,
LISA CHAMBERLAND⁸, JAMES STARRETT⁸, REN-CHUNG CHENG⁹, JONATHAN A. CODDINGTON³,
INGI AGNARSSON^{3,5,10} AND JASON E. BOND⁸

¹Department of Organisms and Ecosystems Research, National Institute of Biology, Večna pot 111, SI-1000, Ljubljana, Slovenia

²Jovan Hadži Institute of Biology, Research Centre of the Slovenian Academy of Sciences and Arts, Novi trg 2, SI-1001, Ljubljana, Slovenia

³Department of Entomology, National Museum of Natural History, Smithsonian Institution, 10th and Constitution, NW, Washington, DC 20560-0105, USA

⁴University of Ljubljana, National Institute of Biology, Večna pot 111, SI-1000, Ljubljana, Slovenia

⁵State Key Laboratory of Biocatalysis and Enzyme Engineering and Centre for Behavioural Ecology and Evolution, School of Life Sciences, Hubei University, Wuhan 430062 Hubei, China

⁶Department of Biotechnology and Systems Biology, National Institute of Biology, Večna pot 111, SI-1000, Ljubljana, Slovenia

⁷Department of Entomology, Plant Pathology and Nematology, University of Idaho, 875 Perimeter Dr. MS 2329, Moscow, ID 83844-2329, USA

⁸Department of Entomology and Nematology, University of California Davis, 1 Shields Ave., Davis, CA 95616, USA

⁹Department of Life Sciences, National Chung Hsing University, No.145 Xingda Rd., South Dist., Taichung City 402, Taiwan and

¹⁰Faculty of Life- and Environmental Sciences, University of Iceland, Sturlugata 7, 102 Reykjavik, Iceland

*Correspondence to be sent to: E-mail: matjaz.kuntner@nib.si.

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Abstract.—Higher-level classifications often must account for monotypic taxa representing depauperate evolutionary lineages and lacking synapomorphies of their better-known, well-defined sister clades. In a ranked (Linnean) or unranked (phylogenetic) classification system, discovering such a depauperate taxon does not necessarily invalidate the rank classification of sister clades. Named higher taxa must be monophyletic to be phylogenetically valid. Ranked taxa above the species level should also maximize information content, diagnosability, and utility (e.g., in biodiversity conservation). In spider classification, families are the highest rank that is systematically catalogued, and *incertae sedis* is not allowed. Consequently, it is important that family-level taxa be well defined and informative. We revisit the classification problem of Orbipurae, an unranked suprafamilial clade containing the spider families Nephilidae, Phonognathidae, and Araneidae *sensu stricto*. We argue that, to maximize diagnosability, information content, conservation utility, and practical taxonomic considerations, this “splitting” scheme is superior to its recently proposed alternative, which lumps these families together as Araneidae *sensu lato*. We propose to redefine Araneidae and recognize a monogeneric spider family, Paraplectanoididae fam. nov. to accommodate the depauperate lineage Paraplectanoides. We present new subgenomic data to stabilize Orbipurae topology which also supports our proposed family-level classification. Our example from spiders demonstrates why classifications must be able to accommodate depauperate evolutionary lineages, for example, Paraplectanoides. Finally, although clade age should not be a criterion to determine rank, other things being equal, comparable ages of similarly ranked taxa do benefit comparative biology. [Classification, family rank, phylogenomics, systematics, monophyly, spider phylogeny.]

In phylogenetic systematics, classifications often need to account for the possibility that monotypic taxa—representing depauperate evolutionary lineages—are discovered as sister to well-defined, ostensibly more diverse, crown groups. This biological reality is due to asymmetric phylogenies and therefore, discovering such a new taxon (that lacks synapomorphies of its better-known sister clade) does not invalidate that clade nor its rank. For example, discovery of a monotypic lineage sister to mammals that lacks mammalian synapomorphies would not invalidate Mammalia.

The Tree of Life is rich with examples of monotypic representatives being sister clades to more speciose ones. For example, *Ginkgo biloba* is the sole representative of the family Ginkgoaceae and order Ginkgoales, and this lineage is sister to Cycadaceae and Cycadales, a clade that contains over a hundred species (Wu et al. 2013). The split between *Ginkgo* and cycads is estimated to have occurred

during the late Carboniferous (Condamine et al. 2015). Similarly, the tuatara (*Sphenodon punctatus*) is the only representative of the family Sphenodontidae and the order Rhynchocephalia, and is sister to Squamata, which includes all remaining lizards and snakes. Here, the split between the clades is estimated to have occurred in the early Mesozoic (Gemmell et al. 2020). Other examples of depauperate evolutionary lineages are *Welwitschia*, *Limmognathia maerski*, and many more.

When depauperate lineages are discovered, the classical Linnean, ranked classification scheme requires naming of all intervening ranks, a so-called Gregg’s paradox (Buck and Hull 1966). A fully rankless higher taxonomic level classification scheme avoids this problem as it continues to accept well-defined clade names even in the light of new, sister lineage discoveries. The International Code of Phylogenetic Nomenclature, for example, does not mandate the use of ranks (Cantino and De Queiroz 2020). A

sensible solution is the combination of the Linnean and phylogenetic systems (Kuntner and Agnarsson 2006), with a limited use of ranks of maximal information content (e.g. genera, families, orders) in a network of phylogenetically defined, unranked clades.

Whether ranked or unranked, named higher taxa must be monophyletic to be phylogenetically valid. Ranked taxa (i.e., above the species level) should also maximize information content, diagnosability, and utility (e.g., communication, comparative biology, cohesive natural history, preservation of traditional concepts, conservation prioritization, etc.). If many well-established depauperate lineages were to be classified into more inclusive taxa (e.g., if Ginkgoaceae were lumped into Cycadaceae), much phylogenetic diversity would be concealed, lowering the taxonomic, conservation, and practical utility of such classification.

EXAMPLE FROM SPIDER CLASSIFICATION

The circumscription of the spider family Araneidae has been debatable, even though the “classic” or “core” common orb weavers are widely and intuitively recognized as a natural group (as well as being the basonym of the order Araneae). The last comprehensive araneid classification was Simon (1895)—taxonomists have been amending it ever since. Araneidae continues to be difficult to diagnose due to the heterogeneity of its contents. Its history is overwhelmingly one of subtracting monophyletic groups to better circumscribe Araneidae, for example, Arkyidae, Cyatholipidae, Linyphiidae, Mimetidae, Nesticidae, Tetragnathidae, and Theridiosomatidae.

Because of downstream implications, the delimitation of family rank taxa in arachnology is more than a semantic issue. In the World Spider Catalog (WSC 2023), a vital, decisive, and universally used tool for arachnologists, the family rank is the terminal rank in which taxa are organized; *incertae sedis* is not considered. Other commonly used family group ranks, such as superfamily, subfamily, and tribe can be found only in the scattered primary literature. For arachnologists, therefore, it is particularly important that family-level taxa be well-defined and informative. Fifteen of the currently recognized spider families are monogeneric (11.3%).

Kuntner et al. (2019) published a phylogenomic study of the spider family Nephilidae recognizing seven genera that included 84% of the known species in the family. This phylogeny, along with earlier generic-level treatments that formed the basis for the removal of the group from Tetragnathidae (Kuntner 2005, 2006, 2007; Harvey et al. 2007; Kuntner et al. 2013), confirmed its monophyly and provided a differential diagnosis with respect to Araneidae, and therefore sharpened the definition of the latter (Kuntner et al. 2019). The argument for the 2019 classification was based on the criteria of monophyly, information content, diagnos-

ability, and in part, inferred clade ages. Kuntner et al. (2019) also recognized Phonognathidae, thus three families: Nephilidae, Phonognathidae, and Araneidae, s.s. Recent phylogenies estimated by other researchers (Dimitrov et al. 2017; Kulkarni et al. 2020; Scharff et al. 2020) agreed that these three clades are mutually monophyletic, well-defined, and informative. Kuntner et al. (2019) named this unranked group of three families Orbipurae, the archetypal orb web spiders, preserving the family group name Araneidae for a classical, core set of diverse, coherent, widely recognized, and most studied groups of genera, including *Araneus*, the type genus of Araneae.

Kallal et al. (2020) revisited the findings of Kallal and Hormiga (2018), which presented a weakly supported topology not confirmed by any subsequent study. Their topology placed Nephilidae and *Paraplectanoides* inside Araneidae *sensu stricto*. Kallal et al.'s (2020) taxonomy creates an expanded Araneidae, which lumps three important, diagnosable families into one, rendering “Araneidae” ambiguous. Although Kallal et al. (2020) address the subfamilies Phonognathinae and Nephilinae, they ignore the implication of these actions, Araneidae *sensu stricto*, providing neither name, diagnosis, nor composition (we acknowledge these were not required). Kallal et al.'s (2020) Araneidae *sensu lato* is equivalent in composition to Orbipurae, but as a family, Araneidae s.l. is challenging to diagnose and loses information content and utility, for example, by uniting into a single family distinct taxa with disparate morphologies and biologies. Such lumping can have consequences. For example, Nephilidae is the only spider family for which all species have been scored for IUCN threat status (<https://www.iucnredlist.org/>) and retaining it as a family adds value to conservation biology. Further, in practice, this results in the loss of classical araneids, phonognathids, and nephilids as entities in WSC and the inability to track these important lineages as such. According to Kallal et al. (2020), a novel placement of the enigmatic Australian *Paraplectanoides* as sister to Nephilidae provides further arguments against Orbipurae, and Nephilidae as a family “renders the family Araneidae paraphyletic.”

Briefly recapped, Kallal et al.'s (2020) arguments for Araneidae s.l. are i) topological: *Paraplectanoides* + Nephilidae nested within Araneidae (and the misplacement of this clade within Araneidae s.s.) negates the family status of Nephilidae; ii) historical: because *Paraplectanoides* has been traditionally an araneid, nephilids are also to be classified as araneids; iii) utilitarian: lumping nontraditional families into a classical one despite diagnosability difficulties and profuse variation in biology simplifies the classification of Araneoidea. Here we use subgenomic data and extensive taxon sampling of Orbipurae to confirm *Paraplectanoides* + Nephilidae, however, that clade is placed outside Araneidae s.s. (as in Kulkarni et al. 2020; Scharff et al. 2020). We then argue that a better supported phylogeny, and the criteria of diagnosability, information content, and utility, eclipse

the arguments put forth by Kallal et al. (2020). We show that the inclusiveness of a family-level taxon directly impacts the utility, information content, and stability of a group's systematic structure. Even if portrayed as arbitrary, decisions whether to lump or split in higher-level classification have consequences that need careful consideration.

Phylogenetic Placement of *Paraplectanoides*

Kallal et al.'s (2020) hypothesized relationship of *Paraplectanoides* + Nephilidae restated results from earlier Sanger data-based hypotheses (Kallal and Hormiga 2018, 2019), placing the pair inside Araneidae s.s. Developing UCE data probes for spiders, Kulkarni et al. (2020) recovered the grouping *Paraplectanoides* + *Trichonephila* outside Araneidae s.s., but the data set contained only one nephilid terminal on a long branch. As the *Paraplectanoides* + *Trichonephila* relationship might plausibly have been an artifact of long-branch attraction (Felsenstein 1978; Bergsten 2005), and as *Paraplectanoides* was not used in the nephilid phylogeny (Kuntner et al. 2019), further testing of this relationship is important. We therefore combined data from the UCE matrix from Kulkarni et al. (2020) with additional UCE data from five nephilid genera, five araneid s.s. genera (selected to maximize subfamily coverage), and two phonognathid genera (See [Supplementary material](#) available on Dryad for methodological details and [Supplementary Table S1](#) for taxon sampling). The strength of this analysis is the inclusion of all but one nephilid genus (*Indoetra*) rather than a select few (Kallal et al. 2020), or a single nephilid representative as in Kulkarni et al. (2020).

Our results (Fig. 1; [Supplementary Figs S1–S4](#) available on Dryad) also recover the sister-group relationship *Paraplectanoides* + Nephilidae and corroborate that detail of the topology of Kallal et al. (2020) and Kulkarni et al. (2020). Unlike Kallal and Hormiga (2018, 2019) and Kallal et al. (2020), our results (and those of Dimitrov et al. 2017; Kulkarni et al. 2020; Scharff et al. 2020) unambiguously place this clade outside Araneidae s.s. The *Cerostris* placement in Kallal and Hormiga (2018, 2019), reused in Kallal et al. (2020), is not supported.

Orbipurae (Phonognathidae ((*Paraplectanoides*, Nephilidae) Araneidae)) is strongly supported in maximum likelihood (Fig. 1; [Supplementary Fig. S2](#) available on Dryad) and coalescent-based (ASTRAL) analyses ([Supplementary Fig. S3](#) available on Dryad). Gene concordance factors (gCF) and site concordance factors (sCF) are positively correlated and are generally high (gCF > 10%; sCF > 40%; Minh et al. 2020) when branch support is high ([Supplementary Figs S3–S5](#) available on Dryad). The tree topology tests support the unconstrained tree over the three alternative tree topologies ([Supplementary Table S2](#) available on Dryad). The relationships within nephilids match the results from Anchored Hybrid Enrichment data (Kuntner et al. 2019) except in the varying position of *Clitaetra*, probably due to low loci recovery, and the absence of *Indoetra* in the current analysis.

To Lump or to Split?

Kallal et al. (2020) state that the Kuntner et al. (2019) classification rendered Araneidae paraphyletic. For various reasons, the taxon sample of Kuntner et al. (2019) did not include *Paraplectanoides*, and Kallal et al.'s (2020) argument was based in part on the weakly supported incorrect placement of Nephilidae + *Paraplectanoides*. Second, the placement of *Paraplectanoides* in their analyses did not *per se* reject the monophyly of any groups we discussed. Therefore, the purported paraphyly was mischaracterized, and as discussed below, can be debated based on topological and logical arguments.

Lumping Nephilidae and Phonognathidae into Araneidae s.l. (Dimitrov et al. 2017; Scharff et al. 2020) reduces the information content of the classification. Kallal et al. (2020) reduced core araneids to the currently nameless, undiagnosed remainder of Araneidae—containing many of the most prominent genera of spiders—unrecognized and untraceable as a group in the World Spider Catalog (WSC 2023). We favor the relimitation of Araneidae to its core taxa and propose an unranked name for the four well-characterized families in Orbipurae. It thus contains one more component of information (Mickevich and Platnick 1989) making an additional statement that Araneidae s.l. does not. At 170 genera, Araneidae s.s. still contains disparate groups and could perhaps be improved in the future by the removal of even more groups, as has happened repeatedly over the last century or more (Scharff et al. 2020).

Kallal et al. (2020) dismiss the possibility of a monogeneric family ranked group for *Paraplectanoides* because that would create a “textbook example of Gregg’s Paradox.” However, Hormiga (1993, 1994) argued against the importance of Gregg’s paradox in the case of Pimoidae, another family that started as monogeneric; similarly, Bond et al. (2020) argued that “nomenclatural stability and abandoning well-established diagnosable genera in favor of a more inclusive polymorphic taxon” favored recognizing the monotypic genus *Cryptocteniza*.

The sister-group *Paraplectanoides* + Nephilidae illustrates Gregg’s paradox. Considering the Linnean requirement that all taxa are unequivocally classified in families, these two spider groups could be classified in different ways. One possibility is to recognize two families (one currently monogeneric). It is possible that other extant or extinct paraplectanoidids will be discovered, as has happened, for example, in Pimoidae, Macrothelidae, Liphistiidae, and Cithaeronidae. *Paraplectanoididae*, rejected by Kallal et al. (2020), allows for stability of the clearly monophyletic and diagnosable Nephilidae, and at the same time, reflects the separate evolutionary history of the two clades. If *Paraplectanoides*, nephilids, and phonognathids are all lumped in Araneidae s.l., the family contains a huge diversity of taxonomic concepts and forms, about which fewer generalizations can be made. As a ranked family, it has lower information content.

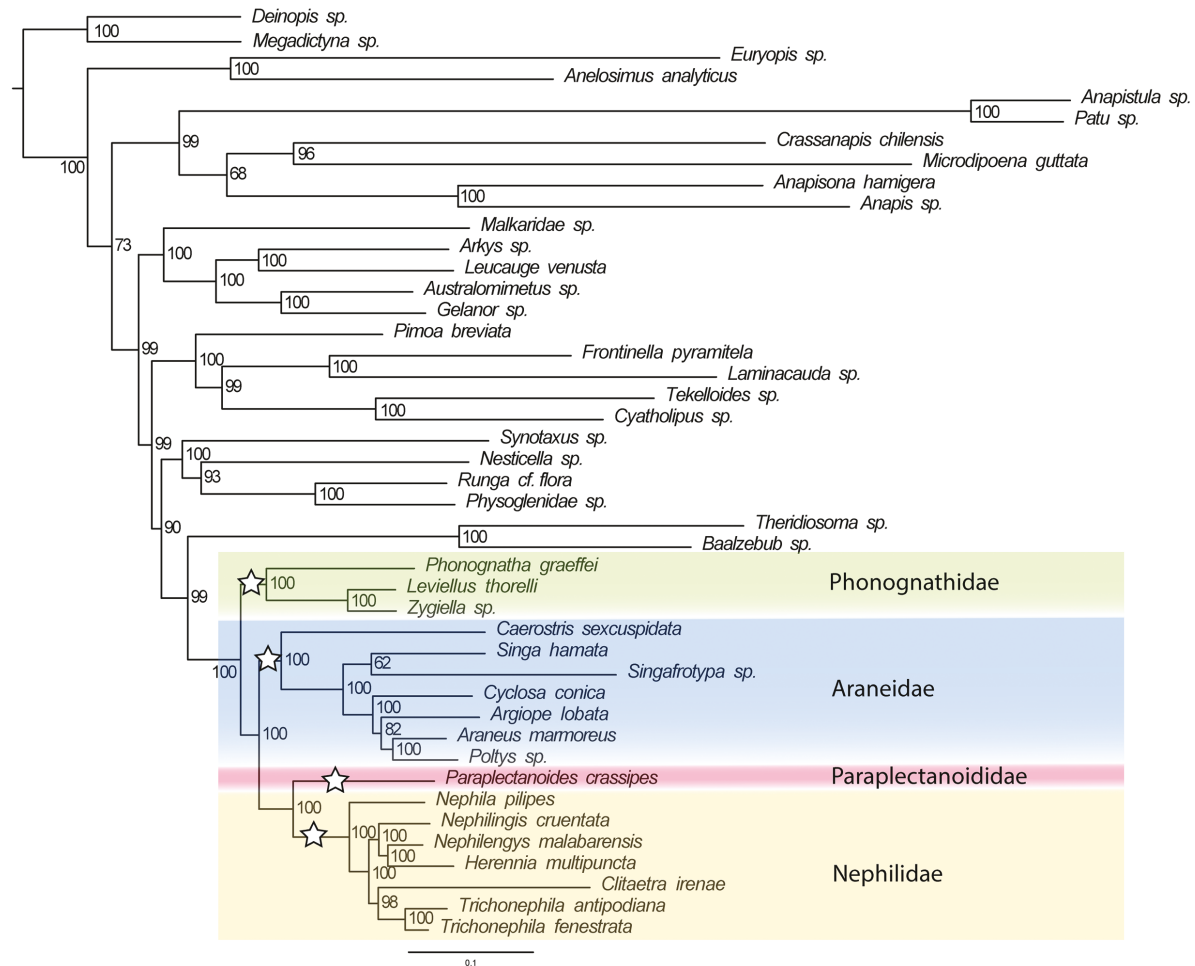


FIGURE 1. The maximum likelihood tree inferred from UCE data for Orbipurae, containing Phonognathidae, Paraplectanoididae, Nephilidae, and Araneidae. Orbipurae families are marked with white stars and their contents colored. Node values represent bootstrap support.

We reexamined *Paraplectanoides* and confirmed that it does not share the key nephilid synapomorphies (see Taxonomy), and thus the first alternative (to recognize two families for *Paraplectanoides* + Nephilidae) better reflects the evolutionary history and diagnosability of each group and maximizes the information content of ranked clades. This avoids the paraphyly of Araneidae and we formally propose this nomenclature in the Taxonomy section below.

Logical Considerations

Rather than redefine Araneidae *s.l.*, Kallal et al. (2020) allege its paraphyly if defined *s.s.* This logic is subject to modal scope fallacy. Modal scope fallacy means that a proposition is taken as necessarily true or false even when the necessity is unwarranted (Schurz 1994; Swartz 2022). Kallal et al.'s (2020) logic seems to be that *Paraplectanoides*, because it is currently catalogued as araneid, will always be an araneid. Therefore, anything related to it must also be an araneid, a modal scope fallacy.

If every genus that is placed as sister to another family retained its prior family affiliation, the family classification of spiders, representing distinct patterns of morphological diversity, would be severely diminished. For example, if another araneid genus is placed phylogenetically as sister to another family, their logic would lump every intervening group within an ever-growing Araneidae *s.l.* At the extreme, all spiders, Araneae, could become araneids, because new sister-group relationships will continue to appear as phylogenetic systematics advances.

Clade Ages and Higher-Level Classification

Kallal et al. (2020) criticize our use of temporal banding demonstrating what a strict application would mean for araneoid spider family circumscription. We agree that strict uniform temporal banding is an impractical and unfeasible goal across the Tree of Life. We nevertheless think that comparable clade ages, if feasible, can add meaning to Linnean ranks. Equally ranked taxa always invite comparison, albeit sometimes unwarranted. Some modern phylogenetic studies have started

to use ages to align classifications within well-defined clades, for example, in birds and mammals (Chen et al. 2019; Delsuc et al. 2019; Presslee et al. 2019; Harvey et al. 2020). Clade age can thus be an informative piece of the evolutionarily informed classification, however, the spider systematics community has not adopted clade age as a criterion to judge alternative classifications. Spider families are typically older than many other animal families. The tendency to lump more sister clades into families (like Araneidae *s.l.*) will likely further increase their age and is in our view an unnecessary step in the wrong direction.

CONCLUSIONS

Most authors agree that monophyly is paramount and that the assignment of ranks in biological classification is in part subjective. Kuntner et al. (2019) proposed that a suite of criteria be decisive in classification decisions to minimize subjectivity. Because our prior call for enhanced repeatability in higher classification was met with criticism (Kallal et al. 2020), herein we provide a rebuttal of that critique. In our view, the key weaknesses of these criticisms are:

1. Kallal et al. (2020) argue that because *Paraplectanoides* is traditionally classified as an araneid, and because phylogenies suggest that it groups as sister to nephilids, nephilids should be classified as araneids. This is a modal scope fallacy.
2. Discovering a sister taxon to a well-defined clade does not, in itself, invalidate that clade's validity or rank as Kallal et al. (2020) suggest for Nephilidae. Classifications sometimes must account for valid but depauperate evolutionary lineages. Hence, *Paraplectanoides* + Nephilidae does not invalidate Nephilidae as a family group, particularly because *Paraplectanoides* lacks key nephilid synapomorphies. The *Caerostris* placement in Kallal et al. (2020) was simply incorrect, and that genus, likewise, does not affect nephilid nomenclature. *Caerostris* is consistently recovered as sister to the remainder of Araneidae and our classification treats the genus accordingly as an araneid.
3. The Araneidae *s.l.* diagnosis in Kallal et al. (2020) is vague and circumscribes a ranked clade of less information content than our proposal because it does not acknowledge the key diagnostic differences that define Nephilidae, Phonognathidae, and Araneidae, which, if recognized, increase the information content of the classification in general (see Taxonomy for diagnosis of Nephilidae, Phonognathidae, and Araneidae). Although Kallal et al. (2020) agree that Nephilidae and Phonognathidae are diagnosable clades, ironically, they neither named nor diagnosed the far larger Araneidae *s.s.* Scharff et al. (2020) recognized the problem and called Araneidae *s.s.* the ARA clade. In other words, one only gets to use the formal family name Araneidae once, and it is better, more conservative, historically more consistent, and more practical to use it to mean what it has historically meant—core Araneidae (see Taxonomy for diagnosis for Araneidae).
4. Kallal et al. (2020) criticized Kuntner et al. (2019) for using clade age as a criterion (temporal banding). Ideally, rank should correlate with age (Chen et al. 2019; Delsuc et al. 2019; Presslee et al. 2019; Harvey et al. 2020), but there should be additional criteria: monophyly, classification information content, and diagnosability. As one way to add objective meaning to ranks, taking clade ages into account can increase the utility of classifications to comparative biology. Araneidae *s.l.* increases the age of Araneidae, and thus the variance in age of families within Araneae and as compared to other animal families.

Our analysis confirms Orbipurae topology and places Paraplectanoididae as sister to the well-defined and diagnosable Nephilidae (Fig. 1). Therefore, in the Taxonomy, we emend Orbipurae to include the monotypic family Paraplectanoididae fam. nov., and revalidate Nephilidae, Phonognathidae, and Araneidae sensu Kuntner et al. (2019).

Our amended classification of Orbipurae eliminates ambiguity in family classification and presents a utilitarian solution with clearly diagnosable, monophyletic families whose biological meaning is well defined. To account for diagnosability and the needs of biodiversity conservation, decisions about lumping or splitting should strongly favor information content. Consequently, classifications must be able to accommodate depauperate evolutionary lineages; discovering them does not necessarily invalidate (i.e., dictate lumping) the rank classification of sister clades. Finally, clade age should not be entirely dismissed in determining rank because, *ceteris paribus*, comparable ages of similarly ranked taxa will continue to benefit comparative biology. Even if often seen as arbitrary, decisions whether to lump or split have pragmatic consequences that need careful consideration.

TAXONOMY

Orbipurae is a clade that contains the families Nephilidae, Phonognathidae, and Araneidae (Kuntner et al. 2019); *Paraplectanoides* is currently classified as an araneid (WSC 2023), but our examination revealed this classification is not based on shared diagnostic features. Based on our results, we propose Paraplectanoididae as a new family in Orbipurae (Araneoidea). Here, we circumscribe all these families, something previously not offered in any one publication. Phylogenetic definitions follow the PhyloCode (Cantino and De Queiroz 2020).

Prior diagnosis of only one of the following clades, Nephilidae, meets modern standards. Paraplectanoid-

idae fam. nov., naming a depauperate lineage, is equal to the diagnosis of the genus *Paraplectanoides* provided here. Phonognathidae is a relatively small family and is easy to diagnose. Araneidae, on the other hand, is vast and has never been well-diagnosed. Below we offer hypotheses for formal diagnoses of these taxa, which will certainly be improved in the future as more taxa and diagnostic characters emerge, and perhaps as disparate and inconsistent groups are removed from Araneidae.

Clade Orbipuræ Kuntner et al. 2019

We define Orbipuræ at the node sometimes discussed as “Araneidae” *sensu lato* (Dimitrov et al. 2017; Kallal et al. 2020; Scharff et al. 2020; WSC 2023). Because Araneidae *sensu lato* has at times contained all the recognized families within Araneoidea, our definition carefully excludes non-Orbipuræ araneoid families.

Definition.—Orbipuræ is the most inclusive crown clade that contains the common ancestor of *Araneus*, *Phonognatha*, *Paraplectanoides*, and *Nephila*, but not of *Theridiosoma*, *Synotaxus*, *Tetragnatha*, *Anapis*, *Theridion*, *Nesticus*, *Linyphia*, *Arkys*, *Cyatholipus*, *Malkara*, *Mimetis*, and *Physoglenes*. This is a maximum-crown-clade definition (Cantino and De Queiroz 2020).

Reference phylogeny.—Figure 1.

Diagnosis.—Spiders that construct two- or three-dimensional webs that usually contain orb web elements, albeit with numerous modifications or losses. Based on available data, Orbipuræ can be diagnosed by a combination of morphological features, for example, the presence of a cheliceral chilum, book lung covers usually grooved, lateral eyes usually on tubercles, and the male palpal median apophysis sharing a hematodocha with the embolic division.

Composition.—The global clade currently contains 188 genera that are catalogued as “araneids” (WSC 2023).

Family Phonognathidae Simon, 1894 New Rank

Definition.—Phonognathidae is the most inclusive crown clade that contains the common ancestor of *Phonognatha*, but not of *Araneus* and not of *Paraplectanoides* and not of *Nephila*. This is a maximum-crown-clade definition (Cantino and De Queiroz 2020).

Reference phylogeny.—Figure 1.

Diagnosis.—Spiders of the family Phonognathidae construct two-dimensional orb webs whose hub is modified into a retreat, either constructed with silk (*Leviellus*) or by using a rolled leaf (*Phonognatha*, *Artiphex*, *Deliochus*, and *Zygiella*). Unlike in nephilids, the cheliceral condyle (boss) in Phonognathidae is smooth and is not striated. Further diagnostic features include distal grouping of setae on palpal tibia and elongated male palpal femur.

Composition.—The family currently contains the genera *Artiphex* Kallal and Hormiga 2022 (a replacement name for *Artifex* Kallal and Hormiga 2018), *Deliochus* Simon 1894, *Leviellus* Wunderlich 2004, *Phonognatha* Simon 1894, and *Zygiella* F. O. Pickard-Cambridge 1902.

Family Araneidae Clerck, 1757

Comments.—We define Araneidae at the node discussed as Araneidae s.s. (Kuntner et al. 2019) and as “the ARA clade” (Scharff et al. 2020). Araneidae will probably continue to be redefined, as advances in comparative morphology, taxonomy, and phylogenetics occur.

Definition.—Araneidae is the most inclusive crown clade that contains the common ancestor of *Araneus*, but not of *Phonognatha* and not of *Paraplectanoides* and not of *Nephila*. This is a maximum-crown-clade definition (Cantino and De Queiroz 2020).

Reference phylogeny.—Figure 1.

Diagnosis.—Spiders of the family Araneidae usually construct two- or three-dimensional orb webs albeit with numerous modifications or losses. Unlike in nephilids, the cheliceral condyle (boss) in Araneidae is smooth, and is not striated. Araneidae may also be diagnosed by a combination of the squat shape of the male palpal tibia, the relatively globular tegulum, eye pattern with lateral eyes widely separated from the medians, and possibly, the presence of an epigynal scape or its homologs.

Composition.—The global family currently contains 175 genera (WSC 2023).

Family Paraplectanoididae Kuntner, Coddington, Agnarsson & Bond fam. nov.

Definition.—*Paraplectanoididae* is the most inclusive crown clade that contains the common ancestor of *Paraplectanoides*, but not of *Araneus* and not of *Phonognatha* and not of *Nephila*. This is a maximum-crown-clade definition (Cantino and De Queiroz 2020).

Reference phylogeny.—Figure 1.

Diagnosis.—Spiders of the family *Paraplectanoididae* construct a three-dimensional nest (Hickman 1975) whose architecture is unlike the archetypal orb web built by most representatives of Orbipuræ. *Paraplectanoides crassipes* builds an ovoid sheet that lacks sticky lines; the only potentially homologous features with the classical orb web are the radii and a horizontal hub (Eberhard 2020). *Paraplectanoididae* males and females uniquely possess a flange on the cheliceral fang (Supplementary Fig. S6 available on Dryad). Females have an additional flange modification of the cheliceral paturon and a highly elevated head region of the carapace (Davies 1988). Unlike in nephilids, the cheliceral condyle (boss) in *Paraplectanoides* is smooth, and is not striated (Supplementary Fig. S6 available on Dryad).

Composition.—The family currently only contains the genus *Paraplectanoides* Keyserling, 1886 with the type species *P. crassipes* Keyserling, 1886 and *P. kochi* O. Pickard-Cambridge, 1877. *Paraplectanoides* is the type genus for the family, by monotypy. Although more species and genera can plausibly be discovered in Australasia or elsewhere on Gondwanan terrains (see Turk et al. 2020), current knowledge suggests that the family and the genus may in fact be monotypic with *P. crassipes* as the only valid species (V. Framenau in litt.), and with *P. kochi* being misplaced in *Paraplectanoides*.

Comments.—Simon (1895) placed *Paraplectanoides* in the group Anepsieae within Argiopidae, but *Anepsion* is a derived araneid not proximal to *Paraplectanoides* (Scharff et al. 2020). Hickman (1975) described the male and provided observations on life history of *P. crassipes*. Although Davies (1988) considered *Paraplectanoides* to be an “araneine” due to its transverse furrows on the epigastric plates and a male palp with a paramedian apophysis and a radix, the key treated the genus as *incertae sedis*.

Family Nephilidae Simon, 1894 Family Rank Resurrected

Definition.—Nephilidae is the most inclusive crown clade that contains the common ancestor of *Nephila*, but not of *Araneus* and not of *Phonognatha* and not of *Paraplectanoides*. This is a maximum-crown-clade definition (Cantino and De Queiroz 2020).

Reference phylogeny.—Figure 1.

Diagnosis.—An unreversed synapomorphy for Nephilidae is also its defining morphological feature: the striated cheliceral boss in both sexes (Kuntner 2005: Figs 6B, 20A–C). In addition, Kuntner et al. (2008) hypothesized 15 unambiguously optimized synapomorphies, as well as additional ambiguously optimized ones that defined Nephilidae. Nephilid lateral eyes are separate in both sexes and male abdomen is sclerotized. Among behavioral characters are those pertaining to web building behavior (the sticky spiral location using the fourth leg tap, double radius construction, double radius attachment on the frame). Nephilid web architectures are either classically orbiculate and aerial (*Nephila*, *Trichonephila*) or modified into elongate, almost rectangular ladder webs against substrate (*Clitaetra*, *Indoetra*, and *Herennia*), or hybrid that are partially aerial and partially substrate anchored, and are always planar, vertical and sticky (Kuntner et al. 2019). However, unlike in Paraplectanoididae, nephilid web modifications are still recognizable through the orb’s classical, homologous elements: frames, hubs, radii, non-sticky and sticky (gluey) spirals (Hormiga et al. 1995; Kuntner et al. 2008; Blackledge et al. 2011; Eberhard 2020), but with sometimes added signal lines (*Nephilengys*, *Nephilingis*; Kuntner 2007), pseudoradii (*Herennia*; Kuntner 2005), or modified hub reinforcements (*Clitaetra*; Kuntner 2006).

Composition.—Nephilidae contains the genera *Clitaetra* Simon, 1889; *Herennia* Thorell, 1877; *Indoetra* Kuntner

2006; *Nephila* Leach, 1815; *Nephilengys* L. Koch, 1872; *Nephilingis* Kuntner 2013; and *Trichonephila* Dahl, 1911. For currently deemed valid species, see Table 1 in Kuntner et al. (2019); other catalogued species names are likely to be proposed as synonyms of these names.

Comments.—Kuntner (2006) proposed Nephilidae Simon, 1894 at the family rank and defined it as the least inclusive clade containing *Clitaetra*, *Herennia*, *Nephila*, and *Nephilengys*. Nephilid monophyly and exclusivity have subsequently been confirmed using molecular and total evidence phylogenetics (Kuntner et al. 2013) and phylogenomics (Kuntner et al. 2019). However, since the above nephilid genera have been further split to reflect their evolutionary history, age and diagnosability, the family Nephilidae now contains species of the seven genera listed above.

SUPPLEMENTARY MATERIAL

Data available from the Dryad Digital Repository: <https://doi.org/10.5061/dryad.t1g1jw5c>.

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CONFLICT OF INTEREST

The authors declare no conflicts of interest.

REFERENCES

- Bergsten J. 2005. A review of long-branch attraction. *Cladistics* 21:163–193.
- Blackledge T.A., Kuntner M., Agnarsson I. 2011. The form and function of spider orb webs: evolution from silk to ecosystems. *Adv. Insect Phys.* 41:175–262.
- Bond J.E., Hamilton C.A., Godwin R.L., Ledford J.M., Starrett J. 2020. Phylogeny, evolution, and biogeography of the North American trapdoor spider family Euctenizidae (Araneae: Mygalomorphae) and the discovery of a new “endangered living fossil” along California’s central coast. *Insect Syst. Divers.* 4:2.
- Buck R.C., Hull D.L. 1966. The logical structure of the Linnaean hierarchy. *Syst. Zool.* XV:97–111.
- Cantino P.D., De Queiroz K. 2020. *International Code of Phylogenetic Nomenclature (PhyloCode)*. Boca Raton: CRC Press, p. 149.

- Chen L., Qiu Q., Jiang Y., Wang K., Lin Z., Li Z., Bibi F., Yang Y., Wang J., Nie W., Su W., Liu G., Li Q., Fu W., Pan X., Liu C., Yang J., Zhang C., Yin Y., Wang Y., Zhao Y., Zhang C., Wang Z., Qin Y., Liu W., Wang B., Ren Y., Zhang R., Zeng Y., Da Fonseca R.R., Wei B., Li R., Wan W., Zhao R., Zhu W., Wang Y., Duan S., Gao Y., Zhang Y.E., Chen C., Hvilsom C., Epps C.W., Chemnick L.G., Dong Y., Mirarab S., Siegmund H.R., Ryder O.A., Gilbert M.T.P., Lewin H.A., Zhang G., Heller R., Wang W. 2019. Large-scale ruminant genome sequencing provides insights into their evolution and distinct traits. *Science* 364(6446):eaav6202.
- Condamine F.L., Nagalingum N.S., Marshall C.R., Morlon H. 2015. Origin and diversification of living cycads: a cautionary tale on the impact of the branching process prior in Bayesian molecular dating. *BMC Evol. Biol.* 15:1–18.
- Davies V.T. 1988. An illustrated guide to the genera of orb-weaving spiders in Australia. *Mem. Queensl. Mus.* 25:273–332.
- Delsuc F., Kuch M., Gibb G.C., Karpinski E., Hackenberger D., Szpak P., Martínez J.G., Mead J.L., McDonald H.G., MacPhee R.D.E., Billet G., Hautier L., Poinar H.N. 2019. Ancient mitogenomes reveal the evolutionary history and biogeography of sloths. *Curr. Biol.* 29:2031–2042.e6.
- Dimitrov D., Benavides L.R., Arnedo M.A., Giribet G., Griswold C.E., Scharff N., Hormiga G. 2017. Rounding up the usual suspects: a standard target-gene approach for resolving the interfamilial phylogenetic relationships of ecribellate orb-weaving spiders with a new family-rank classification (Araneae, Araneioidea). *Cladistics* 33:221–250.
- Eberhard W.G. 2020. *Spider Webs*. Chicago: The University of Chicago Press, p. 658.
- Felsenstein J. 1978. Cases in which parsimony or compatibility methods will be positively misleading. *Syst. Zool.* 27:401–410.
- Gemmell N.J., Rutherford K., Prost S., Tollis M., Winter D., Macey J.R., Adelson D.L., Suh A., Bertozzi T., Grau J.H., Organ C., Gardner P.P., Muffato M., Patricio M., Billis K., Martin F.J., Flicek P., Petersen B., Kang L., Michalak P., Buckley T.R., Wilson M., Cheng Y., Miller H., Schott R.K., Jordan M.D., Newcomb R.D., Arroyo J.I., Valenzuela N., Hore T.A., Renart J., Peona V., Peart C.R., Warmuth V.M., Zeng L., Kortschak R.D., Raison J.M., Zapata V.V., Wu Z., Santesmasses D., Mariotti M., Guigó R., Rupp S.M., Twort V.G., Dussex N., Taylor H., Abe H., Bond D.M., Paterson J.M., Mulcahy D.G., Gonzalez V.L., Barbieri C.G., DeMeo D.P., Pabinger S., Van Stijn T., Clarke S., Ryder O., Edwards S.V., Salzberg S.L., Anderson L., Nelson N., Stone C., Ngatiwai Trust Board. 2020. The tuatara genome reveals ancient features of amniote evolution. *Nature* 584:403–409.
- Harvey M.G., Bravo G.A., Claramunt S., Cuervo A.M., Derryberry G.E., Battilana J., Seeholzer G.F., Shearer McKay J., O'Meara B.C., Faircloth B.C., Edwards S.V., Pérez-Emán J., Moyle R.G., Sheldon F.H., Aleixo A., Smith B.T., Chesser R.T., Silveira L.F., Cracraft J., Brumfield R.T., Derryberry E.P. 2020. The evolution of a tropical biodiversity hotspot. *Science* 370:1343–1348.
- Harvey M.S., Austin A.D., Adams M. 2007. The systematics and biology of the spider genus *Nephila* (Araneae: Nephilidae) in the Australasian region. *Invertebr. Syst.* 21:407–451.
- Hickman V. 1975. On *Paraplectanoides crassipes* Keyserling (Araneae: Araneidae). *Bull. Brit. Arachnol. Soc.* 3:166–174.
- Hormiga G. 1993. Implications of the phylogeny of Pimoidae for the systematics of linyphiid spiders (Araneae, Araneioidea, Linyphiidae). *Mem. Queensl. Mus.* 33:533–542.
- Hormiga G. 1994. A revision and cladistic analysis of the spider family Pimoidae (Araneioidea: Araneae). *Smithson. Contrib. Zool.* 549:1–104.
- Hormiga G., Eberhard W., Coddington J. 1995. Web-construction behavior in Australian *Phonognatha* and the phylogeny of nephilinae and tetragnathid spiders (Araneae, Tetragnathidae). *Aust. J. Zool.* 43:313–364.
- Kallal R.J., Dimitrov D., Arnedo M.A., Giribet G., Hormiga G. 2020. Monophyly, taxon sampling, and the nature of ranks in the classification of orb-weaving spiders (Araneae: Araneioidea). *Syst. Biol.* 69:401–411.
- Kallal R.J., Hormiga G. 2018. Systematics, phylogeny and biogeography of the Australasian leaf-curling orb-weaving spiders (Araneae: Araneidae: Zygiellinae), with a comparative analysis of retreat evolution. *Zool. J. Linn. Soc.* 184:1055–1141.
- Kallal R.J., Hormiga G. 2019. Evolution of the male palp morphology of the orb-weaver hunting spider *Chorizopes* (Araneae: Araneidae) revisited on a new phylogeny of Araneidae, and description of a third species from Madagascar. *Invertebr. Syst.* 6:473–487.
- Kallal R.J., Hormiga G. 2022. *Artiphex* nom. nov. (Araneae: Araneidae), a new replacement name for *Artifex* Kallal & Hormiga, 2018, preoccupied by *Artifex* Silveira, 2008 (Hymenoptera: Vespidae). *Zootaxa* 5175:394.
- Kulkarni S., Wood H., Lloyd M., Hormiga G. 2020. Spider-specific probe set for ultraconserved elements offers new perspectives on the evolutionary history of spiders (Arachnida, Araneae). *Mol. Ecol. Resour.* 20:185–203.
- Kuntner M. 2005. A revision of *Herennia* (Araneae: Nephilidae: Nephilinae), the Australasian “coin spiders.”. *Invertebr. Syst.* 19:391–436.
- Kuntner M. 2006. Phylogenetic systematics of the Gondwanan nephilid spider lineage Clitaetrinae (Araneae, Nephilidae). *Zool. Scr.* 35:19–62.
- Kuntner M. 2007. A monograph of *Nephilengys*, the pantropical “hermit spiders” (Araneae, Nephilidae, Nephilinae). *Syst. Entomol.* 32:95–135.
- Kuntner M., Agnarsson I. 2006. Are the Linnean and phylogenetic nomenclatural systems combinable? Recommendations for biological nomenclature. *Syst. Biol.* 55:774–784.
- Kuntner M., Arnedo M.A., Trontelj P., Lokovšek T., Agnarsson I. 2013. A molecular phylogeny of nephilid spiders: evolutionary history of a model lineage. *Mol. Phylogenet. Evol.* 69:961–979.
- Kuntner M., Coddington J.A., Hormiga G. 2008. Phylogeny of extant nephilid orb-weaving spiders (Araneae, Nephilidae): testing morphological and ethological homologies. *Cladistics* 24:147–217.
- Kuntner M., Hamilton C.A., Cheng R.-C., Gregorič M., Lupše N., Lokovšek T., Lemmon E.M., Lemmon A.R., Agnarsson I., Coddington J.A., Bond J.E. 2019. Golden orbweavers ignore biological rules: phylogenomic and comparative analyses unravel a complex evolution of sexual size dimorphism. *Syst. Biol.* 68:555–572.
- Mickevich M.F., Platnick N.I. 1989. On the information content of classifications. *Cladistics* 5:33–47.
- Minh B.Q., Hahn M.W., Lanfear R. 2020. New methods to calculate concordance factors for phylogenomic datasets. *Mol. Biol. Evol.* 37:2727–2733.
- Presslee S., Slater G.J., Pujos F., Forasiepi A.M., Fischer R., Molloy K., Mackie M., Olsen J.V., Kramarz A., Taglioretti M., Scaglia F., Lezcano M., Lanata J.L., Southon J., Feranec R., Bloch J., Hajduk A., Martin F.M., Salas Gismondi R., Reguero M., de Muizon C., Greenwood A., Chait B.T., Penkman K., Collins M., MacPhee R.D.E. 2019. Palaeoproteomics resolves sloth relationships. *Nat. Ecol. Evol.* 3:1121–1130.
- Scharff N., Coddington J.A., Blackledge T.A., Agnarsson I., Framenau V.W., Szűts T., Hayashi C.Y., Dimitrov D. 2020. Phylogeny of the orb-weaving spider family Araneidae (Araneae: Araneioidea). *Cladistics* 36:1–21.
- Schurz G. 1994. Admissible versus valid rules: a case study of the modal fallacy. *The Monist* 77:376–388.
- Simon E. 1895. *Histoire naturelle des araignées*. Deuxième édition, tome premier. Paris: Roret.
- Swartz N. 2022. Foreknowledge and free will. The Internet Encyclopedia of Philosophy. Available from <https://iep.utm.edu/foreknow/>. Accessed 1 March 2023.
- Turk E., Čandek K., Kralj-Fišer S., Kuntner M. 2020. Biogeographical history of golden orbweavers: chronology of a global conquest. *J. Biogeogr.* 47:1333–1344.
- WSC. 2023. World Spider Catalog Version 24. Available from <http://wsc.nmbe.ch>. Accessed 1 March 2023.
- Wu C.S., Chaw S.M., Huang Y.Y. 2013. Chloroplast phylogenomics indicates that *Ginkgo biloba* is sister to cycads. *Genome Biol. Evol.* 5:243–254.