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# On the formation and the isomer specific detection of methylacetylene ( $\text{CH}_3\text{CCH}$ ), propene ( $\text{CH}_3\text{CHCH}_2$ ), cyclopropane ( $\text{c-C}_3\text{H}_6$ ), vinylacetylene ( $\text{CH}_2\text{CHCCH}$ ), and 1,3-butadiene ( $\text{CH}_2\text{CHCHCH}_2$ ) from interstellar methane ice analogues†

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Pure methane ( $\text{CH}_4$ ) ices processed by energetic electrons under ultra-high vacuum conditions to simulate secondary electrons formed *via* galactic cosmic rays (GCRs) penetrating interstellar ice mantles have been shown to produce an array of complex hydrocarbons with the general formulae:  $\text{C}_n\text{H}_{2n+2}$  ( $n = 4-8$ ),  $\text{C}_n\text{H}_{2n}$  ( $n = 3-9$ ),  $\text{C}_n\text{H}_{2n-2}$  ( $n = 3-9$ ),  $\text{C}_n\text{H}_{2n-4}$  ( $n = 4-9$ ), and  $\text{C}_n\text{H}_{2n-6}$  ( $n = 6-7$ ). By monitoring the *in situ* chemical evolution of the ice combined with temperature programmed desorption (TPD) studies and tunable single photon ionization coupled to a reflectron time-of-flight mass spectrometer, specific isomers of  $\text{C}_3\text{H}_4$ ,  $\text{C}_3\text{H}_6$ ,  $\text{C}_4\text{H}_4$ , and  $\text{C}_4\text{H}_6$  were probed. These experiments confirmed the synthesis of methylacetylene ( $\text{CH}_3\text{CCH}$ ), propene ( $\text{CH}_3\text{CHCH}_2$ ), cyclopropane ( $\text{c-C}_3\text{H}_6$ ), vinylacetylene ( $\text{CH}_2\text{CHCCH}$ ), 1-butyne ( $\text{HCCC}_2\text{H}_5$ ), 2-butyne ( $\text{CH}_3\text{CCCH}_3$ ), 1,2-butadiene ( $\text{H}_2\text{CCCH}(\text{CH}_3)$ ), and 1,3-butadiene ( $\text{CH}_2\text{CHCHCH}_2$ ) with yields of  $2.17 \pm 0.95 \times 10^{-4}$ ,  $3.7 \pm 1.5 \times 10^{-3}$ ,  $1.23 \pm 0.77 \times 10^{-4}$ ,  $1.28 \pm 0.65 \times 10^{-4}$ ,  $4.01 \pm 1.98 \times 10^{-5}$ ,  $1.97 \pm 0.98 \times 10^{-4}$ ,  $1.90 \pm 0.84 \times 10^{-5}$ , and  $1.41 \pm 0.72 \times 10^{-4}$  molecules  $\text{eV}^{-1}$ , respectively. Mechanistic studies exploring the formation routes of methylacetylene, propene, and vinylacetylene were also conducted, and revealed the additional formation of the 1,2,3-butatriene isomer. Several of the above isomers, methylacetylene, propene, vinylacetylene, and 1,3-butadiene, have repeatedly been shown to be important precursors in the formation of polycyclic aromatic hydrocarbons (PAHs), but until now their interstellar synthesis has remained elusive.

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## 1. Introduction

During the last few decades, methylacetylene ( $\text{CH}_3\text{CCH}$ ), propene ( $\text{CH}_3\text{CHCH}_2$ ), vinylacetylene ( $\text{CH}_2\text{CHCCH}$ ), and 1,3-butadiene ( $\text{CH}_2\text{CHCHCH}_2$ ) have received considerable attention from the astrochemistry and astronomy communities due to their critical roles in astrochemical processes.<sup>1,2</sup> Interestingly, each of these molecules has been shown to be capable of forming precursors to polycyclic aromatic hydrocarbons (PAHs), prototype PAHs, and PAHs themselves.<sup>3</sup> Specifically, methylacetylene emerged as a precursor in the formation of PAHs such as indene ( $\text{C}_9\text{H}_8$ )

as well as 5- and 6-methyl-1*H*-indene ( $\text{C}_{10}\text{H}_{10}$ ) through the bimolecular reaction with a phenyl radical ( $\text{C}_6\text{H}_5$ ) and *via* a reaction with the *para*-tolyl radical ( $\text{C}_6\text{H}_4\text{CH}_3$ ), respectively.<sup>4-7</sup> Methylacetylene has also been used in models to explain Titan's atmospheric chemistry.<sup>8-11</sup> Similarly, propene ( $\text{CH}_3\text{CHCH}_2$ ) produced resonantly stabilized free radicals (RSFRs) – precursors to PAHs – in gas-phase reactions with carbon atoms ( $\text{C}$ ) and dicarbon molecules ( $\text{C}_2$ ) forming the methylpropargyl radical ( $\text{C}_4\text{H}_5$ ) as well as 1- and 3-vinylpropargyl radical ( $\text{C}_5\text{H}_5$ ), respectively.<sup>12-14</sup> Also, vinylacetylene ( $\text{CH}_2\text{CHCCH}$ ) has been suggested as a hydrocarbon important for Titan's chemistry,<sup>11,15,16</sup> and forms prototype PAHs such as naphthalene ( $\text{C}_{10}\text{H}_8$ ), 1- and 2-methylnaphthalene ( $\text{C}_{11}\text{H}_{10}$ ), and *ortho*-benzynes ( $\text{o-C}_6\text{H}_4$ ) in the gas phase.<sup>17-20</sup> Finally, 1,3-butadiene ( $\text{CH}_2\text{CHCHCH}_2$ ) is also fundamental as an essential building block in the gas-phase growth of prototypical PAHs through reactions with dicarbon ( $\text{C}_2$ ) to form the phenyl radical ( $\text{C}_6\text{H}_5$ ), with the tolyl radical ( $\text{C}_6\text{H}_4\text{CH}_3$ ) forming 6-methyl-1,4-dihydronaphthalene ( $\text{C}_{11}\text{H}_{12}$ ), and with the ethynyl radical ( $\text{CCH}$ ) producing benzene ( $\text{C}_6\text{H}_6$ ).<sup>21-23</sup>

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The  $C_3H_4$  isomer methylacetylene ( $CH_3CCH$ ) was first observed towards Sagittarius B2 and Orion A *via* the  $5_0 \rightarrow 4_0$  transitions at 85 GHz.<sup>24,25</sup> Subsequently, methylacetylene has been detected in several astronomical surveys toward TMC-1, SgrB2, PKS 1830-211, L1544, NGC 7538 IRS9, tentatively in NCG 4418, and recently for the first time in a photo-dissociation region *via* the Horsehead nebula at fractional abundances – in all sources – of a few  $10^{-9}$  with respect to molecular hydrogen.<sup>26–33</sup> This results in a column density of about  $8 \times 10^{13}$  molecules  $cm^{-2}$ .<sup>34</sup> Besides the interstellar medium (ISM), the methylacetylene molecule has also been observed in our Solar System on Jupiter, Saturn, Uranus, and Titan.<sup>35–38</sup> Many years later the  $C_3H_6$  isomer propene ( $CH_3CHCH_2$ ) was detected in the ISM using 13 rotational lines from 84 to 104 GHz toward TMC-1 and more recently in the cold cores Lupus-1A, L1495B, L1521F, and Serpens South 1a with fractional abundances of a few  $10^{-9}$ .<sup>34,39</sup> This calculates to a column density of about  $4 \times 10^{13}$  molecules  $cm^{-2}$ .<sup>34</sup> Propene has also been detected in the atmosphere of Titan.<sup>40</sup> However, none of the other  $C_3$  hydrocarbons, allene ( $CH_2CCH_2$ ), cyclopropene ( $c-C_3H_4$ ), or cyclopropane ( $c-C_3H_6$ ), has been detected in the ISM. Although the  $C_4$  hydrocarbons vinylacetylene ( $H_2CCHCCH$ ) and 1,3-butadiene ( $H_2CCHCHCH_2$ ) have remained elusive in the ISM they are still astrophysically important molecules. Vinylacetylene has been shown to react with the phenyl radical and produce the PAH naphthalene<sup>1,18</sup> and has been suggested to be an important hydrocarbon for Titan's chemistry.<sup>11,15,16</sup> Similarly, 1,3-butadiene can react with the ethynyl radical ( $CCH$ ) to form benzene ( $C_6H_6$ ),<sup>23</sup> with the phenyl radical ( $C_6H_5$ ) to form 1,4-dihydronaphthalene ( $C_{10}H_{10}$ ),<sup>41</sup> or with the tolyl radical ( $C_7H_7$ ) to form 5- and 6-methyl-1,4-dihydronaphthalene ( $C_{11}H_{12}$ ).<sup>42</sup>

Recently, it was shown that the methylacetylene molecule cannot be synthesized *via* gas-phase reactions alone.<sup>27,43</sup> An extensive review of the chemical pathways leading to methylacetylene in the gas phase concluded that there are no efficient gas-phase synthetic pathways available to reproduce the observed abundances of this molecule in the cold ISM.<sup>27</sup> Here, the main gas-phase source of methylacetylene was proposed to be through ion-molecule reactions that produce  $C_3H_5^+$  followed by dissociative recombination reactions with minor formation pathways *via* neutral-neutral reactions. Since the gas-phase reactions were not able to reproduce the ISM abundance of methylacetylene, surface reactions taking place on interstellar grains have been investigated. These gas-phase species were assumed to stick to the cold dust grains and then undergo hydrogenation at an ultra-cold ISM temperature of 10 K. The methylacetylene molecule was shown to form *via* the hydrogenation of  $C_3$  which could then be further hydrogenated to propene. However, the model including gas-phase and grain-surface reactions cannot reproduce the observed methylacetylene abundance by more than an order of magnitude. These failed attempts to reproduce the methylacetylene abundance show that important formation pathways are still missing at low temperatures.<sup>27,28,33</sup>

It has also been proposed that the propene molecule was synthesized *via* gas-phase reactions. However, a comprehensive analysis of these hypothesized reactions shows that some have considerable entrance barriers, which makes them inefficient

in the cold ISM.<sup>27,44</sup> Following the failure to reproduce the observations of propene in the ISM with pure gas-phase mechanisms surface reactions on low temperature ISM grains were considered as an alternate pathway once again. Similar to methylacetylene, models incorporating solely the surface reactions producing propene – from the hydrogenation of methylacetylene – fail to reproduce the observed abundance by an order of magnitude.<sup>27</sup> Another pathway investigated the recombination of radicals, from exploding ice mantles, in a high-density gas-phase model, but this again failed to replicate the observed abundances.<sup>45–47</sup> Although these models show a better agreement than pure gas-phase models to the observed abundance there is clearly a lack of a significant production route at ISM temperatures.<sup>27,34</sup>

Although, vinylacetylene and 1,3-butadiene have remained undetected in the ISM, several reactions have shown that they can be produced under ISM conditions. Vinylacetylene has been shown to be a product of several gas-phase reactions such as the reaction of the ethynyl radical with ethylene ( $C_2H_4$ ),<sup>11,48,49</sup> the ethynyl radical with propene,<sup>48</sup> and the methylidyne radical ( $CH$ ) with propene.<sup>50</sup> Vinylacetylene has also been a proposed product in several theoretical gas-phase studies, some modeling Titan's atmosphere, from the reaction of the ethynyl radical with ethylene ( $C_2H_4$ ), propene ( $C_3H_6$ ), and 1-butene ( $C_4H_8$ ).<sup>51,52</sup> The formation of vinylacetylene *via* solid-phase pathways has also been hypothesized such as in acetylene ( $C_2H_2$ ) ices *via* the recombination of a vinyl radical ( $C_2H_3$ ) and an ethynyl radical,<sup>53</sup> or possibly through the recombination of excited acetylene molecules.<sup>54</sup> Utilizing some of these reaction pathways vinylacetylene has been incorporated into ISM chemical models.<sup>55–57</sup> Also, 1,3-butadiene can be formed in gas-phase reactions such as the reaction of propene with the methylidyne radical,<sup>23,50,58,59</sup> as well as the reaction of the methylidyne radical with *trans*-butene.<sup>50</sup> This important molecule has also been speculated to form in ices through the recombination of two vinyl radicals.<sup>23</sup> Although multiple studies have been carried out on the production pathways of these four molecules – ethylacetylene, propene, vinylacetylene, 1,3-butadiene – their formation pathways remain mostly speculative. The lack of agreement of models, using these reactions, with observation shows that they are ill-defined and clearly there are missing synthetic routes producing these astrochemically important molecules.

Here, we demonstrate that methylacetylene, propene, vinylacetylene, and 1,3-butadiene can be formed in pure methane ice when exposed to ionizing radiation in the form of energetic electrons which mimic secondary electrons produced from the passage of galactic cosmic rays (GCRs) through ISM ices.<sup>60–63</sup> Although no pure methane ice has been detected in space it is an ice constituent at levels up to 11% with respect to water in the ISM<sup>64</sup> and as a major constituent on Pluto *via* the recent New Horizons mission.<sup>65,66</sup> The current study utilizes single photon ionization coupled with a reflectron time-of-flight mass spectrometer (PI-ReTOF-MS) to detect methylacetylene, propene, vinylacetylene, and the 1,3-butadiene isomers selectively. Traditionally, Fourier-transform infrared spectroscopy (FTIR) and electron impact quadrupole mass spectrometry (EI-QMS)

have been used to identify molecules formed in ISM analog ices that have been exposed to ionizing radiation. However, FTIR can only firmly identify individual molecules that are very small, such as carbon monoxide (CO), water (H<sub>2</sub>O), carbon dioxide (CO<sub>2</sub>), and methane (CH<sub>4</sub>), all of which have been detected as constituents of ISM ices, or functional groups of more complex molecules.<sup>64</sup> As molecules become more complex their infrared vibrations begin to overlap and make definitive identification based solely on FTIR analysis impossible. Fig. 1 shows an overlay of mid-IR spectra of the hydrocarbons methane (CH<sub>4</sub>), ethylene (C<sub>2</sub>H<sub>4</sub>), ethane (C<sub>2</sub>H<sub>6</sub>), allene (CH<sub>2</sub>CCH<sub>2</sub>), methylacetylene (CH<sub>3</sub>CCH), propane (C<sub>3</sub>H<sub>8</sub>), and butane (C<sub>4</sub>H<sub>10</sub>), which clearly show that the hydrocarbons larger than C<sub>2</sub> begin to have extensive overlap of their most intense vibrations making their identification very difficult. Gas-phase analysis of the subliming molecules from the irradiated ice can provide complementary assignments to FTIR analysis. Typically this is accomplished with EI-QMS, but this technique often utilizes an ionization energy of 70 eV which not only ionizes most organic molecules but also causes their extensive fragmentation. Fig. 2 shows an overlay of the fragmentation pattern of the hydrocarbons methane (CH<sub>4</sub>), ethylene (C<sub>2</sub>H<sub>4</sub>), ethane (C<sub>2</sub>H<sub>6</sub>), allene (CH<sub>2</sub>CCH<sub>2</sub>), methylacetylene (CH<sub>3</sub>CCH), propane (C<sub>3</sub>H<sub>8</sub>), and butane (C<sub>4</sub>H<sub>10</sub>). Here, several of the molecules' molecular ions are not the most intense signal corresponding to the parent molecule and many of the fragments of these hydrocarbons have overlapping masses. In order to provide a more sensitive analysis of the processed ice, these limited capabilities of FTIR and EI-QMS have been further complemented by incorporating single photon ionization coupled with a reflectron time-of-flight mass spectrometer (PI-ReTOF-MS), which allows isomer specific analysis of the subliming products

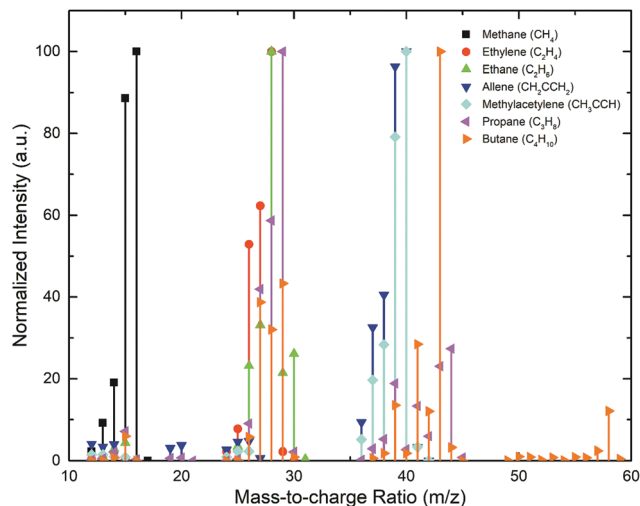


Fig. 2 Overlay of electron ionization, at 70 eV, fragmentation patterns of several hydrocarbons (<https://webbook.nist.gov/chemistry/>).

from the ISM analog ices.<sup>67–76</sup> This technique utilizes soft photon ionization which results in minimal or no fragmentation of the parent molecule.<sup>68,69,77</sup> Furthermore, the tunable photoionization energy allows for specific isomers to be identified based on each isomer's ionization energy.<sup>78–83</sup>

## 2. Experimental

All experiments were completed in an ultrahigh vacuum (UHV) chamber at pressures of typically  $3 \times 10^{-11}$  Torr.<sup>68,84,85</sup> A UHV compatible closed-cycle helium compressor (Sumitomo Heavy

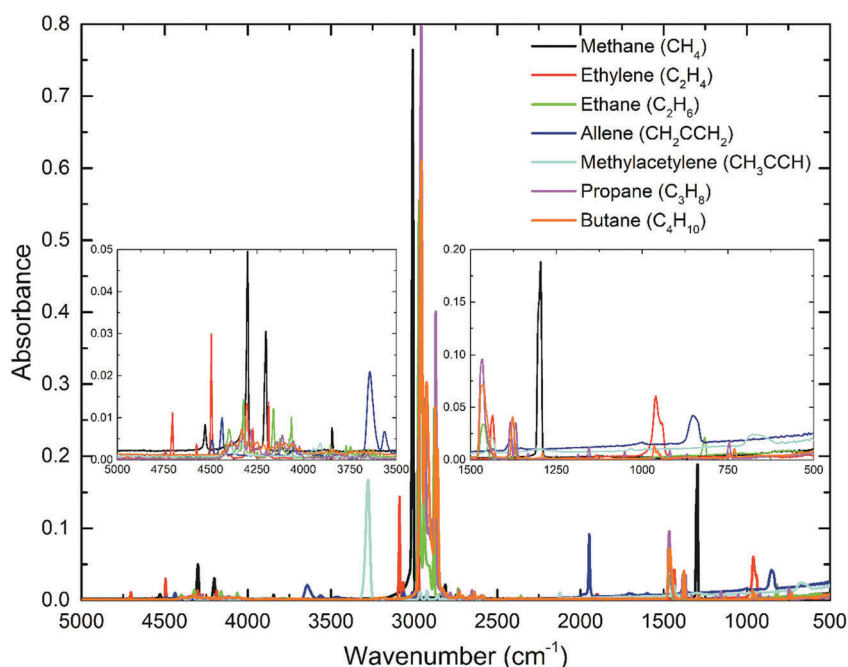


Fig. 1 Infrared spectra at 5000–500 cm<sup>-1</sup> of several pure unprocessed ices at 5 K. The left inset is a 16× magnification of 5000–3500 cm<sup>-1</sup> and the right inset is a 4× magnification of 1500–500 cm<sup>-1</sup>.

Industries, RDK-415E) connected to a cold finger, constructed with oxygen free high conductivity copper, is interfaced to the UHV chamber using a differentially pumped rotary feedthrough (Thermionics Vacuum Products, RNN-600/FA/MCO) and a UHV compatible bellow (McAllister, BLT106) that allows it to be rotated in the horizontal plane and/or translated in the vertical plane, respectively. The substrate, a highly reflective silver mirror, is interfaced to the cold finger *via* indium foil to ensure thermal conductivity. Also, a glass capillary array is used to deposit the gases onto the silver target. Typical background pressures of  $5 \times 10^{-8}$  Torr in the UHV chamber over a few minutes resulted in the desired thickness of methane (Specialty Gases of America, 99.999%) ice to be deposited.<sup>86</sup> During this deposition the ice thickness was determined by monitoring on line and *in situ* the interference pattern produced during the gas deposition *via* a HeNe laser ( $\lambda = 632.8$  nm; CVI Melles-Griot; 25-LHP-230) reflecting off of the silver substrate into a photodiode.<sup>87–89</sup> The ice thickness was determined to be  $590 \pm 50$  nm from this method by using a refractive index  $n = 1.280 \pm 0.008$ .<sup>90</sup> Using a modified Lambert–Beer relationship with absorption coefficients of  $3.95 \times 10^{-19}$ ,  $1.40 \times 10^{-17}$ ,  $1.29 \times 10^{-20}$ ,  $3.89 \times 10^{-19}$ ,  $8.15 \times 10^{-19}$ , and  $8.76 \times 10^{-20}$  cm molecule<sup>-1</sup> and integrated areas for the respective infrared bands 2814 ( $\nu_2 + \nu_4$ ), 3010 ( $\nu_3$ ), 4114 ( $\nu_2 + 2\nu_4$ ), 4202 ( $\nu_1 + \nu_4$ ), 4301 ( $\nu_3 + \nu_4$ ), and 4528 ( $\nu_2 + \nu_3$ ) yielded an average thickness of  $440 \pm 130$  nm, which is in agreement with the more accurate laser interferometry method.<sup>90</sup>

Once the ice was deposited a Fourier-transform infrared spectrometer (FTIR; Nicolet 6700), which probes the active infrared vibrational modes of the reactant and product molecules within the sample, was utilized to continuously analyze the ice on line and *in situ* before, during, and after processing. The FTIR was operated in absorption–reflection–absorption mode at a reflection angle of  $45^\circ$  to the substrate normal in the mid-IR region from 5000 to 500 cm<sup>-1</sup> with a resolution of 4 cm<sup>-1</sup> at 2 minute intervals, which allowed collection of 30 spectra during the one hour of irradiation with 5 keV electrons.<sup>91–94</sup> Following the irradiation phase the ice was held isothermal at 5 K for one hour. Next, the substrate was heated to 300 K through a controlled ramp rate of 0.5 K min<sup>-1</sup> (temperature programmed desorption; TPD). The electron gun (specs EQ 22-35) is positioned 100 cm from the sample, and the energetic electrons processed  $1.0 \pm 0.1$  cm<sup>2</sup> of the methane ice striking it at an incidence angle of  $70^\circ$  relative to the surface normal of the substrate for 1 hour with a current of 30 nA, which results in a fluence of  $(6.7 \pm 0.5) \times 10^{14}$  electrons cm<sup>-2</sup>.<sup>95,96</sup> Utilizing Monte Carlo simulations *via* CASINO 2.42 software<sup>97</sup> the mean average penetration depth of the energetic electrons was calculated to be  $410 \pm 20$  nm and only up to 5% of the energy is absorbed by the ice beyond 250 nm. Using a density of 0.47 g cm<sup>-3</sup> for methane it was determined that the average energy deposited into the ice was  $3.5 \pm 1.1$  eV molecule<sup>-1</sup> of methane (Table S1, ESI†).<sup>98–100</sup> It was found that ices inside cold dense clouds in the ISM receive an energy dose of 0.3–3 eV molecule<sup>-1</sup> over  $10^6$ – $10^7$  years from GCRs, which is the energetic processing that is attempted to be replicated in the present manuscript. Furthermore, although

the low flux of interstellar like photons and GCRs can be achieved no experimental laboratory can exactly mimic ISM irradiation conditions due to the length of the residence times of the ices ( $>10^6$  years), and therefore the total dose absorbed by the ice is a reasonable way to compare the processing of ices across experimental labs and to the ISM.<sup>101</sup> Also, as seen from the average penetration depth of the electrons, chemical reactions take place throughout the bulk of the ice which has traditionally been considered inert and not considered in astrochemical models.<sup>79</sup>

During the TPD phase, the subliming molecules were monitored *via* a QMS (Extrel, Model 5221) and PI-ReTOF-MS. Here, the QMS instrument monitors a mass range from 1–300 amu with a 70 eV electron impact ionization source and an emission current of 2 mA by operating in a residual gas analyzer (RGA) mode.<sup>102–104</sup> In order to monitor the subliming molecules with PI-ReTOF-MS the substrate was rotated  $180^\circ$  to face the ReTOF and cannot be monitored by FTIR at the same time, thus requiring multiple experiments for each system. The PI-ReTOF-MS technique involves ionizing subliming molecules *via* single photon ionization in the form of pulsed coherent VUV light.<sup>105–107</sup> The generation of the VUV light used was accomplished *via* four-wave difference mixing, which requires the interaction of three photons in a non-linear medium to produce a fourth VUV photon. Here, the photons generated from a Nd:YAG laser (Spectra-Physics, PRO-270-30 or PRO-250-30) were used directly to produce VUV at an energy of 10.49 eV (118 nm), *via* non-resonant four-wave mixing, or acted as a pump laser for a separate dye laser. Two dye lasers were used that have the capability of tuning from 350–630 nm (Sirah, Cobra-Stretch) or 400–900 nm (Sirah, Precision Scan), and either dye laser output can be frequency doubled, or the former dye laser output can be frequency tripled with BBO crystals. The laser outputs, from the Nd:YAG and/or dye lasers, are then focused into a pulse valve housing emitting the non-linear medium, which was either krypton or xenon in the present experiments. After the interaction region containing the non-linear medium a lithium fluoride (LiF) lens is used to separate the input wavelengths from the desired VUV wavelength utilizing the difference in refractive indexes across wavelengths for LiF. The VUV light was then directed in front of the substrate to ionize the subliming molecules, which were then detected with the ReTOF. Finally, the intensity was then measured with a NIST calibrated photodiode, which is typically optimized for each VUV energy to around  $10^{12}$  photons s<sup>-1</sup>. To distinguish which C<sub>3</sub>H<sub>4</sub>, C<sub>3</sub>H<sub>6</sub>, C<sub>4</sub>H<sub>4</sub>, and C<sub>4</sub>H<sub>6</sub> isomers were formed in the processed methane ice four different photoionization energies of 10.49 eV ( $\lambda = 118.2$  nm), 9.77 eV ( $\lambda = 126.9$  nm), 9.45 eV ( $\lambda = 131.2$  nm), and 9.15 eV ( $\lambda = 135.5$  nm) were employed (Fig. 3). Next, once a molecule was ionized it was detected utilizing a reflectron time-of-flight mass spectrometer (Jordan TOF products, Inc.). Here, the ions were detected *via* a dual chevron configured multichannel plate (MCP). The MCP signal was then amplified with a fast pre-amplifier (Ortec 9305) followed by shaping *via* a 100 MHz discriminator. A personal-computer-based multichannel scaler (FAST ComTec, P7888-1 E) then recorded the



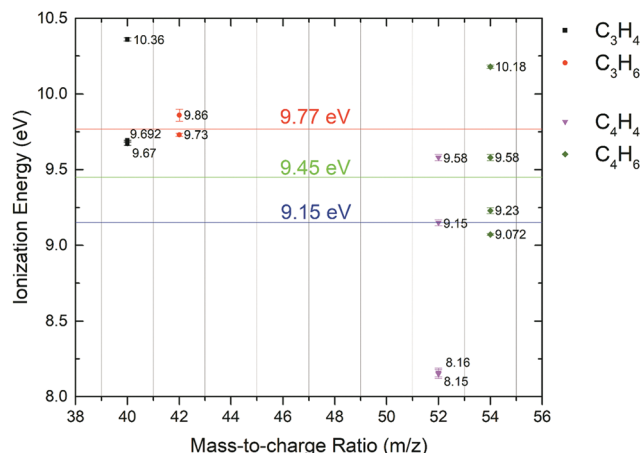


Fig. 3 Scheme for PI energy to discriminate the  $C_3H_4$  isomers (black squares),  $C_3H_6$  isomers (red circles),  $C_4H_4$  isomers (pink triangles), and  $C_4H_6$  isomers (green diamonds).

spectra using 4 ns bin widths triggered at 30 Hz (Quantum Composers, 9518), with 3600 sweeps in a mass spectrum which corresponds to a 1 K change in temperature of the substrate per mass spectrum.

### 3. Results

#### 3.1 Infrared spectroscopy

A complete analysis utilizing the FTIR spectrum of the electron irradiated methane ice has been previously discussed.<sup>100</sup> Briefly, six products propane ( $C_3H_8$ ), ethane ( $C_2H_6$ ), ethyl radical ( $C_2H_5$ ), ethylene ( $C_2H_4$ ), acetylene ( $C_2H_2$ ), and the methyl radical ( $CH_3$ ) were detected *via* several infrared absorptions.<sup>108,109</sup> Kinetic fitting of the temporal profiles *via* the numerical solution of a system of coupled differential equations allowed the extraction of a reaction mechanism producing the small molecules observed in the infrared spectrum and was extrapolated to explain the more complex species observed *via* PI-ReTOF-MS.<sup>100</sup> However, Fig. 4 shows infrared stretches that can be assigned to several small

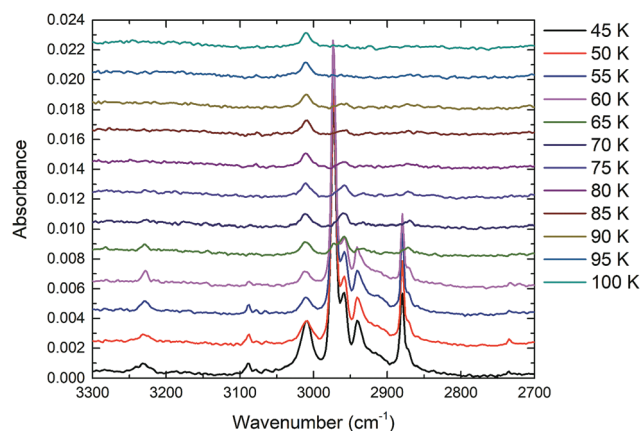


Fig. 4 Infrared spectra at 3300–2700  $cm^{-1}$  after methane ice irradiation during TPD after the majority of the methane reactant has sublimed (see the text for peak assignments).

hydrocarbons propane ( $C_3H_8$ ,  $\nu_s$ , 2961  $cm^{-1}$ ), ethane ( $C_2H_6$ ,  $\nu_s$ , 2883  $cm^{-1}$ ), ethylene ( $C_2H_4$ ,  $\nu_9$ , 3093  $cm^{-1}$ ), acetylene ( $C_2H_2$ ,  $\nu_3$ , 3266  $cm^{-1}$ ), and methane ( $CH_4$ ,  $\nu_3$ , 3008  $cm^{-1}$ ) and yet several of these infrared stretches remain well past the known sublimation temperature of their assignments, showing that they have contributions to their signal from other more complex species with higher sublimation temperatures. By comparing literature values or depositing pure samples of methylacetylene, propene, vinylacetylene, and 1,3-butadiene and recording their infrared spectrum it is clear that several of these infrared stretches belong to these molecules (ESI†; Fig. S1; Tables S2 and S3). Furthermore, by monitoring these infrared stretches during TPD several can be correlated with sublimation events of methylacetylene *via* its  $\nu_6$  fundamental at 2959  $cm^{-1}$  and propene by the  $\nu_3$  (2970  $cm^{-1}$ ),  $\nu_{15}$  (2940  $cm^{-1}$ ),  $\nu_4$  (2915  $cm^{-1}$ ), and  $2\nu_7$  (2883  $cm^{-1}$ ) infrared peaks (Fig. S2, ESI†). However, many more complex hydrocarbons could also be detected with PI-ReTOF-MS.

#### 3.2 PI-ReTOF-MS

During TPD the subliming molecules were monitored *via* PI-ReTOF-MS (PI = 10.49 eV; 118 nm), which detected products belonging to five hydrocarbon groups with the generic formulae:  $C_nH_{2n+2}$  ( $n = 4-8$ ),  $C_nH_{2n}$  ( $n = 3-9$ ),  $C_nH_{2n-2}$  ( $n = 3-9$ ),  $C_nH_{2n-4}$  ( $n = 4-9$ ), and  $C_nH_{2n-6}$  ( $n = 6-7$ ) (Fig. 5). Interestingly, ions corresponding to the molecular formulae  $C_3H_4$ ,  $C_3H_6$ ,  $C_4H_4$ , and  $C_4H_6$  were all detected in this experiment which could correspond to the astrophysically important molecules methylacetylene ( $m/z = 40$ ), propene ( $m/z = 42$ ), vinylacetylene ( $m/z = 52$ ), and 1,3-butadiene ( $m/z = 54$ ), respectively (Fig. 6). Each of the above molecular formulae has multiple isomers associated with them and therefore the contribution due to each isomer to the overall signal is unknown. Therefore, several experiments utilizing tunable photoionization were needed to further discriminate which specific isomers were formed from the electron irradiated methane ice (ESI†, Table S4).

**3.2.1  $C_3H_4$ .** The ion signal detected at  $m/z = 40$  can only correspond to the molecular formula  $C_3H_4$ , which has three possible isomers: methylacetylene ( $CH_3CCH$ ; IE =  $10.36 \pm 0.01$  eV), allene ( $CH_2CCH_2$ ; IE =  $9.69 \pm 0.01$  eV), and cyclopropene ( $c-C_3H_4$ ; IE =  $9.67 \pm 0.01$  eV) (Fig. 6a).<sup>110</sup> The ion signal began to be detected at about 75 K, peaked at 87 K, and continued to sublime up to 110 K where it then returned to the baseline. To better determine the identity of these signals calibration experiments were also performed (ESI†). Here, a mixture containing 1% methylacetylene, 1% propene, and 1% 1,3-butadiene in methane was deposited and then monitored during TPD with PI-ReTOF-MS (PI = 10.49 eV) (Fig. S3, ESI†). This calibration experiment detected a similar peak at  $m/z = 40$  subliming from 75–110 K. Despite the very close match of the unknown ion signal to the known methylacetylene signal it could still be due to any of the three isomers, but most likely had some contribution from the methylacetylene isomer. In order to definitively identify the isomers contributing to the signal the PI energy was tuned to 9.77 eV (Fig. 6e). Interestingly, no signal was detected at  $m/z = 40$  at this PI energy, and therefore it is only due to the

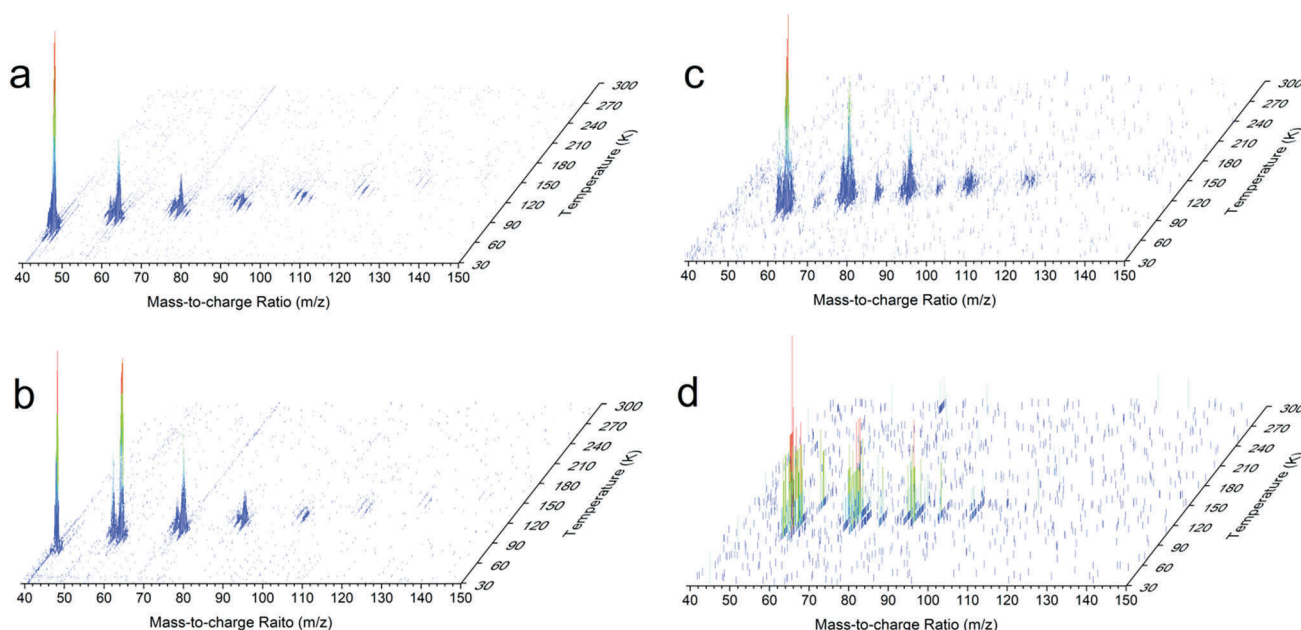


Fig. 5 PI-ReTOF-MS data versus temperature at PI energies of (a) 10.49 eV, (b) 9.77 eV, (c) 9.45 eV, and (d) 9.15 eV. The signal intensity is color coded where blue corresponds to 1–25%, cyan to 25–50%, green to 50–75%, and red to 75–100% of the max intensity detected in each graph.

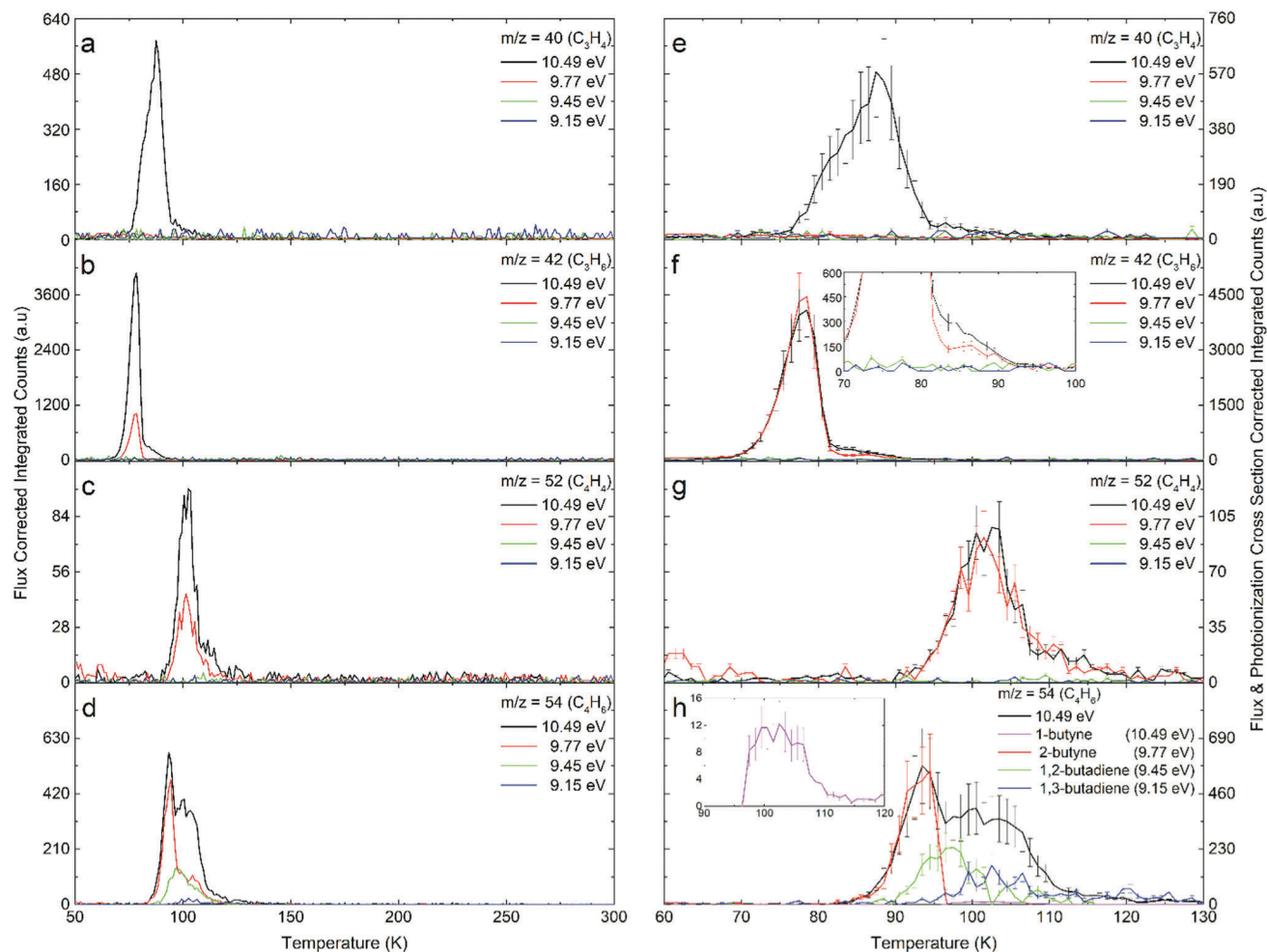
methylacetylene isomer and neither of the other two isomers. As expected no ion signal was detected at PI energies of 9.45 eV or 9.15 eV either. Furthermore, by accounting for the photoionization cross section at 10.49 eV of 18.9 Mb and utilizing the PI-ReTOF-MS correction factor derived from the calibration ices the yield of methylacetylene was calculated to be  $2.17 \pm 0.95 \times 10^{-4}$  molecules  $\text{eV}^{-1}$  (Table 1 and Table S4, ESI†).

**3.2.2  $\text{C}_3\text{H}_6$ .** The signal detected at  $m/z = 42$  is due to a hydrocarbon with the molecular formula  $\text{C}_3\text{H}_6$ , which can only be due to cyclopropane ( $\text{c-C}_3\text{H}_6$ ; IE =  $9.86 \pm 0.04$ ) or propene ( $\text{CH}_3\text{CHCH}_2$ ; IE =  $9.73 \pm 0.01$ ).<sup>110</sup> Here, the signal ranged from 68–91 K and peaked at 77 K (Fig. 6b). The calibration experiment signal of propene showed a similar onset sublimation temperature of 67 K (Fig. S3, ESI†). Although the onset sublimation temperature of propene matches nicely with the unknown  $m/z = 42$  signal it could also be due to cyclopropane as either isomer can be ionized at a PI of 10.49 eV. Switching to a PI energy of 9.77 eV, the signal is still detectable at  $m/z = 42$ , which can only be due to propene (Fig. 6f). This detection unambiguously identifies propene as a contributor to the ion signal. However, further investigation of the ion signals – after accounting for differences in flux of the experiments and the photoionization cross section of the isomers (Table S4, ESI†) – detected with PI energies of 10.49 eV and 9.77 eV reveals that a small portion of the 10.49 eV signal, where both isomers are able to be ionized, from about 80–90 K was due to the cyclopropane isomer (Fig. 6f inset). As expected no ion signal was detected at PI energies of 9.45 eV or 9.15 eV because neither of these isomers will be photoionized. After applying the experimentally derived correction factor from the calibration ices to the photoionization cross section corrected data the yields of propene and cyclopropane were calculated to be  $3.7 \pm 1.5 \times 10^{-3}$  molecules  $\text{eV}^{-1}$

and  $1.23 \pm 0.77 \times 10^{-4}$  molecules  $\text{eV}^{-1}$ , respectively (Table 1). This shows a much larger production of propene and corresponds to a propene to cyclopropane ratio of  $30 \pm 22 : 1 \pm 0.75$ .

**3.2.3  $\text{C}_4\text{H}_4$ .** A signal was also detected at  $m/z = 52$ , which corresponds to the molecular formula  $\text{C}_4\text{H}_4$  and could be due to vinylacetylene ( $\text{CH}_2\text{CHCCH}$ ; IE =  $9.58 \pm 0.02$ ), 1,2,3-butatriene ( $\text{CH}_2\text{CCCH}_2$ ; IE =  $9.16 \pm 0.02$ ), cyclobutadiene ( $\text{c-C}_4\text{H}_4$ ; IE =  $8.16 \pm 0.03$ ), and/or methylenecyclopropene ( $\text{CH}_2(\text{c-C}_3\text{H}_2)$ ; IE =  $8.15 \pm 0.03$ ) (Fig. 6c).<sup>110</sup> The sublimation profile of  $m/z = 52$  extended from 94–120 K. Tuning the PI energy to 9.77 eV revealed no change in the ion signal, within the error bars, after correcting for differences in photon flux and isomer cross sections (Fig. 6g). This result is expected as all of the above isomers are able to be photoionized at both 10.49 eV and 9.77 eV. However, adjusting the PI energy to 9.45 eV resulted in no signal detection at  $m/z = 52$  and therefore the only  $\text{C}_4\text{H}_4$  isomer produced in the methane ice can be unmistakably assigned to vinylacetylene (Fig. 6g). Also, no signal was detected at PI = 9.15 eV. Following the previously described procedure the yield of vinylacetylene produced from the irradiated methane ice was calculated to be  $1.90 \pm 0.84 \times 10^{-5}$  molecules  $\text{eV}^{-1}$  (Table 1).

**3.2.4  $\text{C}_4\text{H}_6$ .** PI-ReTOF-MS also detected an ion signal at  $m/z = 54$  belonging to the molecular formula  $\text{C}_4\text{H}_6$ , which may be due to any of the four different isomers: 1,3-butadiene ( $\text{H}_2\text{CCHCHCH}_2$ ; IE = 9.07 eV), 1,2-butadiene ( $\text{H}_2\text{CCCH}(\text{CH}_3)$ ; IE = 9.03 eV), 1-butyne ( $\text{HCCC}_2\text{H}_5$ ; IE = 10.18 eV), and/or 2-butyne ( $\text{CH}_3\text{CCCH}_3$ ; IE = 9.58 eV) (Fig. 6d).<sup>110</sup> The ion signal was observed to sublime from 85–115 K with multiple peak events, which may be due to several different isomers subliming at different temperatures. However, the previously described calibration sublimation profile of 1,3-butadiene, subliming from 87–115 K, matched very nicely with the unknown ion



**Fig. 6** PI-ReTOF-MS photon flux corrected signals for (a)  $m/z = 40$  ( $C_3H_4$ ), (b)  $m/z = 42$  ( $C_3H_6$ ), (c)  $m/z = 52$  ( $C_4H_4$ ), (d)  $m/z = 54$  ( $C_4H_6$ ), as well as photoionization cross section corrected signals (e)  $m/z = 40$  ( $C_3H_4$ ), (f)  $m/z = 42$  ( $C_3H_6$ ), (g)  $m/z = 52$  ( $C_4H_4$ ), (h)  $m/z = 54$  ( $C_4H_6$ ) at PI energies of 10.49 eV (black), 9.77 eV (red), 9.45 eV (green), and 9.15 eV (blue). Panel h profiles correspond to the noted isomer in the legend determined via subtraction of overlapping isomers (see the text). The green (9.45 eV) and blue (9.15 eV) traces have been magnified by a factor of 10 in panels a and e, and a factor of 20 in panels b and f to be visible.

**Table 1** Yields of detected  $C_3H_4$ ,  $C_3H_6$ ,  $C_4H_4$ , and  $C_4H_6$  isomers

Molecule	Molecules $eV^{-1}$	Relative ratio
$C_3H_4$ Methylacetylene	$2.17 \pm 0.95 \times 10^{-4}$	—
$C_3H_6$ Propene	$3.7 \pm 1.5 \times 10^{-3}$	$30 \pm 22 : 1 \pm 0.75$ (propene : cyclopropane)
Cyclopropane	$1.23 \pm 0.77 \times 10^{-4}$	
$C_4H_4$ Vinylacetylene	$1.90 \pm 0.84 \times 10^{-5}$	—
$C_4H_6$ 1,3-Butadiene	$1.41 \pm 0.72 \times 10^{-4}$	$3.5 \pm 2.5 : 4.9 \pm 3.4 : 1 \pm 0.7 : 3.2 \pm 2.3$ (1,3-butadiene : 1,2-butadiene : 2-butyne : 1-butyne)
1,2-Butadiene	$1.97 \pm 0.98 \times 10^{-4}$	
2-Butyne	$4.01 \pm 1.98 \times 10^{-5}$	
1-Butyne	$1.28 \pm 0.65 \times 10^{-4}$	

signal suggesting that the signal is at least partially due to this isomer (Fig. S3, ESI<sup>†</sup>). To discriminate which isomers contributed to  $m/z = 54$  the PI energy was changed to 9.77 eV, 9.45 eV, and 9.15 eV to no longer ionize certain isomers (Fig. 3). At all of the PI energies  $m/z = 54$  was detected, showing that

there could be contributions from multiple isomers (Fig. 6d). However, the detection of  $m/z = 54$  at 9.15 eV unambiguously identifies 1,3-butadiene as a product. After correcting the ion signal, for flux differences and each isomer's photoionization cross sections at each PI, the amount that each isomer contributed to

the total signal at 10.49 eV could be calculated (Table S4, ESI†). First, the signal detected at 9.15 eV can only belong to 1,3-butadiene, and because the signal can be corrected to the amount that it contributed to the signal at 9.45 eV it can be subtracted from the total signal detected for  $m/z = 54$  at 9.45 eV. The remaining signal can then only be due to 1,2-butadiene. This procedure was repeated for the signal at 9.77 eV by subtracting the corrected 1,3-butadiene and 1,2-butadiene signals to determine the contribution from 2-butyne. This procedure was repeated as well for the 10.49 eV ion signal by subtracting the influences of 1,3-butadiene, 1,2-butadiene, and 2-butyne to determine the amount of 1-butyne produced (Fig. 6h). The yields for 1,3-butadiene, 1,2-butadiene, 2-butyne, and 1-butyne were determined to be  $1.41 \pm 0.72 \times 10^{-4}$  molecules  $\text{eV}^{-1}$ ,  $1.97 \pm 0.98 \times 10^{-4}$  molecules  $\text{eV}^{-1}$ ,  $4.01 \pm 1.98 \times 10^{-5}$  molecules  $\text{eV}^{-1}$ , and  $1.28 \pm 0.65 \times 10^{-4}$  molecules  $\text{eV}^{-1}$ , respectively, which corresponds to a ratio of  $3.5 \pm 2.5:4.9 \pm 3.4:1 \pm 0.7:3.2 \pm 2.3$  (Table 1).

## 4. Discussion

### 4.1 Reaction mechanism & energetics

In order to deduce the mechanistic details about the synthesis of methylacetylene, propene, and vinylacetylene a retro-synthesis of each molecule was performed to understand the different pathways, and ultimately what starting compounds to use (Fig. 7). The complexity of the 1,3-butadiene formation pathways is beyond the scope of the current manuscript. The energetics of these pathways are investigated to help explain why certain mechanisms are viable while others remain inaccessible (Table 2). Furthermore, these mechanisms may provide insights into the relative ratios determined above.

**4.1.1  $\text{C}_3\text{H}_4$ .** Methylacetylene could be formed *via* three possible pathways: insertion of carbene ( $\text{CH}_2$ ) into an acetylene carbon-hydrogen bond, recombination of a methyl radical ( $\text{CH}_3$ ) with an ethynyl radical ( $\text{CCH}$ ), and addition of a methyl radical ( $\text{CH}_3$ ) to acetylene forming the allyl radical ( $\text{CH}_3\text{CHCH}$ )

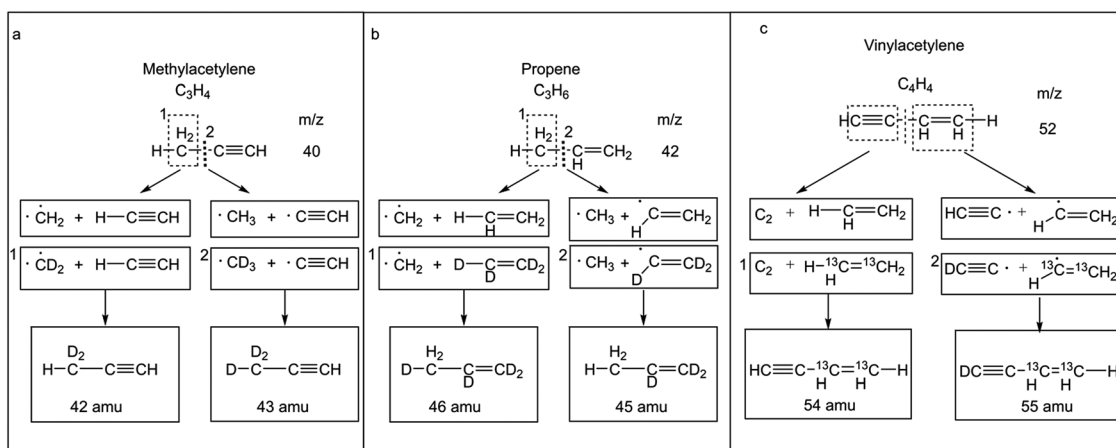


Fig. 7 Mechanisms from isotopic substitution for (a) methylacetylene, (b) propene, and (c) vinylacetylene.

Table 2 Reaction pathways and energetics for methylacetylene, propene, and vinylacetylene

Reaction	Reaction	Reaction energy ( $\Delta_R G$ ) <sup>a</sup>		Reaction barrier ( $E_b$ )	
		$\text{kJ mol}^{-1}$	eV	$\text{kJ mol}^{-1}$	eV
1	$\text{CH}_4(\text{X}^1\text{A}_1) \rightarrow \text{CH}_3(\text{X}^2\text{A}_2'') + \text{H}(\text{X}^2\text{S}_{1/2})$	429	4.44	—	—
2	$\text{CH}_4(\text{X}^1\text{A}_1) \rightarrow \text{CH}_2(\text{a}^1\text{A}_1) + \text{H}_2(\text{X}^1\Sigma_g^+)$	495	5.13	—	—
3	$\text{C}_2\text{H}_2(\text{X}^1\Sigma_g^+) \rightarrow \text{C}_2\text{H}(\text{X}^2\Sigma^+) + \text{H}(\text{X}^2\text{S}_{1/2})$	553	5.73	—	—
4	$\text{CH}_3(\text{X}^2\text{A}_2'') + \text{C}_2\text{H}(\text{X}^2\Sigma^+) \rightarrow \text{CH}_3\text{CCH}(\text{X}^1\text{A}')$	−523	−5.42	—	—
5	$\text{CH}_2(\text{a}^1\text{A}_1) + \text{C}_2\text{H}_2(\text{X}^1\Sigma_g^+) \rightarrow \text{CH}_3\text{CCH}(\text{X}^1\text{A}')$	−465	−4.82	—	—
6	$\text{CH}_3(\text{X}^2\text{A}_2'') + \text{C}_2\text{H}_2(\text{X}^1\Sigma_g^+) \rightarrow \text{CH}_3\text{CCH}(\text{X}^1\text{A}') + \text{H}(\text{X}^2\text{S}_{1/2})$	30	0.31	33 <sup>b</sup>	0.34 <sup>b</sup>
7	$\text{C}_2\text{H}_4(\text{X}^1\text{A}_g) \rightarrow \text{C}_2\text{H}_3(\text{X}^2\text{A}') + \text{H}(\text{X}^2\text{S}_{1/2})$	454	4.71	—	—
8	$\text{CH}_2(\text{a}^1\text{A}_1) + \text{C}_2\text{H}_4(\text{X}^1\text{A}_g) \rightarrow \text{CH}_3\text{CHCH}_2(\text{X}^1\text{A}')$	−455	−4.72	—	—
9	$\text{CH}_3(\text{X}^2\text{A}_2'') + \text{C}_2\text{H}_3(\text{X}^2\text{A}') \rightarrow \text{CH}_3\text{CHCH}_2(\text{X}^1\text{A}')$	−414	−4.29	—	—
10	$\text{CH}_3(\text{X}^2\text{A}_2'') + \text{C}_2\text{H}_4(\text{X}^1\text{A}_g) \rightarrow \text{CH}_3\text{CHCH}_2(\text{X}^1\text{A}') + \text{H}(\text{X}^2\text{S}_{1/2})$	40	0.41	185 <sup>b</sup>	1.92 <sup>b</sup>
11	$\text{C}_2\text{H}(\text{X}^2\Sigma^+) \rightarrow \text{C}_2(\text{X}^1\Sigma_g^+) + \text{H}(\text{X}^2\text{S}_{1/2})$	473	4.90	—	—
12	$\text{C}_2(\text{X}^1\Sigma_g^+) + \text{C}_2\text{H}_4(\text{X}^1\text{A}_g) \rightarrow \text{CH}_2\text{CHCCH}(\text{X}^1\text{A}')$	−588	−6.09	—	—
13	$\text{C}_2\text{H}(\text{X}^2\Sigma^+) + \text{C}_2\text{H}_3(\text{X}^2\text{A}') \rightarrow \text{CH}_2\text{CHCCH}(\text{X}^1\text{A}')$	−569	−5.90	—	—
14	$\text{C}_2\text{H}_2(\text{X}^1\Sigma_g^+) + \text{C}_2\text{H}_3(\text{X}^2\text{A}') \rightarrow \text{CH}_2\text{CHCCH}(\text{X}^1\text{A}') + \text{H}(\text{X}^2\text{S}_{1/2})$	−16	−0.17	6.2 <sup>c</sup>	0.06 <sup>c</sup>
15	$\text{C}_2\text{H}(\text{X}^2\Sigma^+) + \text{C}_2\text{H}_4(\text{X}^1\text{A}_g) \rightarrow \text{CH}_2\text{CHCCH}(\text{X}^1\text{A}') + \text{H}(\text{X}^2\text{S}_{1/2})$	−115	−1.19	— <sup>c</sup>	— <sup>c</sup>

<sup>a</sup> Reaction energies using NIST Chemistry WebBook (<https://webbook.nist.gov/chemistry/>). <sup>b</sup> Ryazantsev *et al.* (2015). <sup>c</sup> Krishtal *et al.* (2009).



followed by hydrogen loss producing methylacetylene (Fig. 7a). In order to determine the methylacetylene reaction mechanism mixed ices of D4-methane ( $\text{CD}_4$ ) and acetylene ( $\text{C}_2\text{H}_2$ ), with a ratio of 1:1, were irradiated to unravel which pathways produced methylacetylene in methane ices (Fig. 7a).<sup>111,112</sup> Here, the detection of  $m/z = 42$  ( $\text{C}_3\text{H}_2\text{D}_2$ ) shows that methylacetylene was produced *via* insertion of the carbene. Alternatively, the detection of  $m/z = 43$  ( $\text{C}_3\text{HD}_3$ ) would prove that the radical recombination of the methyl radical with the ethynyl radical occurs to produce methylacetylene. It should be noted that the third pathway, *via* methyl radical addition to acetylene, results in an identical mass-to-charge ratio as the radical recombination route and therefore these two pathways are indecipherable no matter which isotopic substitution scheme is used.

Fig. 8 shows that both  $m/z = 42$  and 43 were detected subliming from the D4-methane–acetylene ice at a photoionization energy of 10.49 eV. These ion signals could belong to any of the deuterium substituted  $\text{C}_3\text{H}_4$  isotopomers ( $m/z = 42$  and 43) or possibly from  $\text{C}_3\text{H}_6$  isomers ( $m/z = 42$ ). To determine which molecules were formed, several tunable experiments were carried out. Utilizing a PI energy of 9.93 eV will photoionize all  $\text{C}_3\text{H}_4$  isotopomers *except* methylacetylene ( $\text{CH}_3\text{CCH}$ ; IE =  $10.36 \pm 0.01$  eV) and all  $\text{C}_3\text{H}_6$  isomers. Therefore, any change in  $m/z = 42$  or 43 can be correlated with methylacetylene. Here, we see that  $m/z = 43$  drops to the baseline and thus the only molecule responsible for the  $m/z = 43$  signal is methylacetylene (Fig. 8b). Similarly, Fig. 8a displays the change in  $m/z = 42$  at 9.93 eV (red line) from 10.49 eV (black dashed line), and the difference of these two signals can only correspond to methylacetylene (orange line). Here, the methylacetylene (orange line) is the difference spectrum produced by subtracting the 9.93 eV

signal from the 10.49 eV signal. Furthermore, because the signal at  $m/z = 43$  at 10.49 eV was solely due to methylacetylene this peak was scaled and subtracted from  $m/z = 42$  at 10.49 eV, and an overlay of this difference spectrum with the  $m/z = 42$  ion signal recorded at PI = 9.93 eV matches very well confirming the assignment of methylacetylene (Fig. S4a, ESI†).

However, there are still other ions adding to the signal  $m/z = 42$  at 10.49 eV (black dashed line). By changing the PI energy to 9.77 eV we can further constrain what molecules are responsible for this signal as any change from 9.93 eV to 9.77 eV will be due to the contribution of cyclopropane ( $\text{c-C}_3\text{H}_6$ ; IE =  $9.86 \pm 0.04$  eV). At PI = 9.77 eV (blue line) another change in the  $m/z = 42$  TPD profile is observed confirming the presence of cyclopropane, but a signal is still present (Fig. 8a). To determine the cyclopropane contribution (cyan line) the signal recorded for  $m/z = 42$  at 9.77 eV (blue line) was subtracted from the signal recorded at 9.93 eV (red line). The remaining signal at 9.77 eV (blue line) could be due to the  $\text{C}_3\text{H}_4$  isotopomers allene ( $\text{D}_2\text{CCCH}_2$ ; IE =  $9.69 \pm 0.01$  eV), or cyclopropene ( $\text{c-C}_3\text{H}_2\text{D}_2$ ; IE =  $9.67 \pm 0.01$  eV), and/or the  $\text{C}_3\text{H}_6$  isomer propene ( $\text{CH}_3\text{CHCH}_2$ ; IE =  $9.73 \pm 0.01$  eV). However, by irradiating a pure acetylene ice an identical signal at  $m/z = 42$  was detected, which can only be due to  $\text{C}_3\text{H}_6$  isomers, and is most likely the propene isomer due to its match with the 9.77 eV peak at  $m/z = 42$  from the D4-methane–acetylene ice experiment (Fig. S4b, ESI†). Using this information it was possible to determine what portion of the 9.93 eV spectrum was due to cyclopropane (Fig. 8a).

Now that the viable reaction pathways have been detected the reactions' energetics to produce methylacetylene can be discussed (Table 2). The formation of the methyl radical (reaction (1)) and carbene (reaction (2)) is endoergic by 4.44 eV ( $429 \text{ kJ mol}^{-1}$ ) and

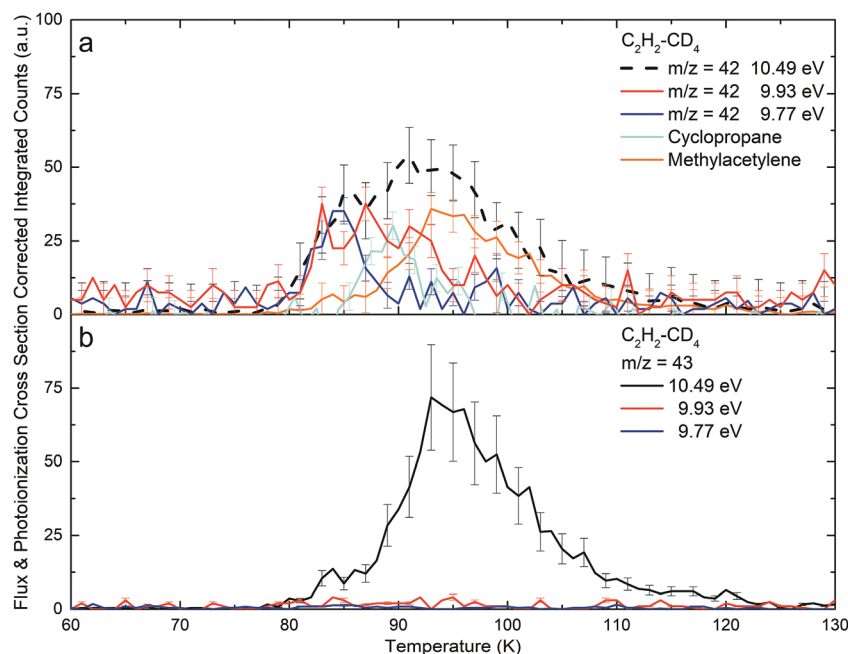


Fig. 8 PI-ReTOF-MS flux and photoionization cross section corrected signals using photoionization energies of 10.49 eV (black), 9.93 eV (red), and 9.77 eV (blue) for (a)  $m/z = 42$  which was determined to be due to methylacetylene (orange), cyclopropane (cyan), and most likely propene (blue) and (b)  $m/z = 43$  from the irradiated  $\text{C}_2\text{H}_2$ – $\text{CD}_4$  ice mixture (see the text for details).

5.13 eV (495 kJ mol<sup>-1</sup>), respectively, which can be accounted for by the impinging electrons (Table S1, ESI†). Also, the formation of the ethynyl radical is endoergic and requires 5.73 eV (553 kJ mol<sup>-1</sup>; reaction (3)) to form. The radical recombination pathway for the formation of methylacetylene *via* the recombination of the methyl and ethynyl radicals has no barrier and is exoergic by -5.42 eV (-523 kJ mol<sup>-1</sup>; reaction (4)). Alternatively, the insertion of the carbene into the acetylene molecule is also barrierless, when in the singlet excited state (<sup>1</sup>A<sub>1</sub>), and exoergic by -4.82 eV (-465 kJ mol<sup>-1</sup>; reaction (5)). However, the final pathway of methyl radical addition to the acetylene molecule (reaction (6)) has a barrier of 0.34 eV (33 kJ mol<sup>-1</sup>) and is endoergic by 0.31 eV (30 kJ mol<sup>-1</sup>).<sup>113,114</sup> The energetics of these pathways show that both the radical recombination (reaction (4)) and insertion (reaction (5)) pathways are feasible as they are barrierless and exoergic, but that the addition route (reaction (6)) is unlikely at the low temperature of the experiment (5 K) under thermal equilibrium as it has a barrier and is also an endoergic reaction. Here, the detection of *m/z* = 42 and 43 with the methylacetylene portion in a ratio of 1.0 ± 0.2 : 2.0 ± 0.4, respectively, shows that the radical recombination pathway to methylacetylene is preferred (reaction (4)). This preference matches with the energetics as the formation of the methyl radical and recombination of it with the ethynyl radical are less endoergic and more exoergic, respectively, when compared to the formation of carbene and its insertion into the acetylene molecule. In summary, methylacetylene is formed *via* radical recombination and carbene insertion in a ratio of 2.0 ± 0.4 : 1.0 ± 0.2.

**4.1.2 C<sub>3</sub>H<sub>6</sub>.** Similarly, propene is able to be formed through three routes: insertion of carbene (CH<sub>2</sub>) into an ethylene

carbon-hydrogen bond, recombination of a methyl radical (CH<sub>3</sub>) with a vinyl radical (C<sub>2</sub>H<sub>3</sub>), and addition of a methyl radical (CH<sub>3</sub>) to ethylene forming the propyl radical (CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>) followed by hydrogen loss forming propene. Propene's reaction mechanism was probed *via* mixed ices of methane and D<sub>4</sub>-ethylene in a ratio of 1 : 1 (Fig. 7b). Detection of *m/z* = 45 from this experiment shows that propene is produced *via* radical recombination of the methyl radical with the vinyl radical. Alternatively, the detection of *m/z* = 46 means that propene was produced from the insertion of the carbene into an ethylene carbon-hydrogen bond. Again, it should be noted that the third pathway, *via* methyl radical addition using the vinyl radical, results in an identical mass-to-charge ratio as the radical recombination route. The results of these experiments are shown in Fig. 9, and confirmed the presence of both *m/z* = 45 and 46 at a photoionization energy of 10.49 eV. Here, the only ions that can be associated with *m/z* = 45 and 46 are the propene and cyclopropane isotopomers. By tuning the photoionization energy to 9.77 eV only propene (IE = 9.73 ± 0.01 eV) will be ionized and not the cyclopropane (IE = 9.86 ± 0.04 eV) isotopomer. The results of this tunable experiment confirm the formation of propene *via* both the radical recombination pathway as well as the insertion route as an ion signal was detected at both *m/z* = 45 and 46. Here, radical recombination formed solely the propene isotopomer (Fig. 9a). However, after accounting for the flux and photoionization cross section of propene at PI = 9.77 eV the *m/z* = 46 TPD profile had a portion of its signal that could not be accounted for by only propene. Therefore, a portion of the *m/z* = 46 signal at 10.49 eV must belong to cyclopropane (Fig. 9b).

As stated above for methylacetylene, the formation of the methyl radical and carbene are endoergic by 4.44 eV (429 kJ mol<sup>-1</sup>) and 5.13 eV (495 kJ mol<sup>-1</sup>), respectively (reactions (1) and (2)).

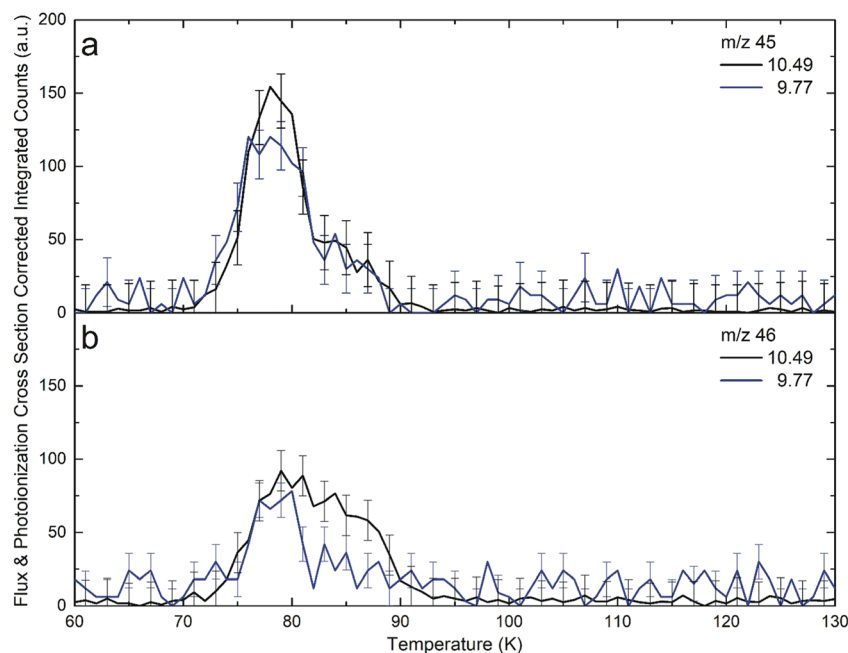


Fig. 9 PI-ReTOF-MS flux and photoionization cross section corrected signals for (a) *m/z* = 45 and (b) *m/z* = 46 using photoionization energies of 10.49 eV (black) and 9.77 eV (blue), from the irradiated C<sub>2</sub>D<sub>4</sub>-CH<sub>4</sub> ice mixture.

Also, the formation of the vinyl radical is endoergic and requires 4.71 eV (454 kJ mol<sup>-1</sup>; reaction (7)) to form. The insertion of the carbene into the ethylene molecule is barrierless, if the carbene is in the singlet excited state ( $a^1A_1$ ), and exoergic by -4.72 eV (-455 kJ mol<sup>-1</sup>; reaction (8)). The recombination of the methyl and ethynyl radicals also has no barrier though and is exoergic by -4.29 eV (-414 kJ mol<sup>-1</sup>; reaction (9)). Finally, the addition of the methyl radical to ethylene leading to propene and a hydrogen atom (reaction (10)) has a barrier of 1.92 eV (185 kJ mol<sup>-1</sup>) and is endoergic by 0.41 eV (40 kJ mol<sup>-1</sup>).<sup>113,114</sup> Again, the addition reaction mechanism involving a barrier is not likely under thermal conditions due to the low temperature of the system. The detection of propene *via*  $m/z = 45$  and 46 in a ratio of  $1.5 \pm 0.4 : 1 \pm 0.25$ , at 9.77 eV, shows that radical recombination is preferred for propene synthesis (reaction (9)). This is again reflected by the energetics as reactions (8) and (9) have very similar exoergicities. However, it is possible that the carbene needed to form propene is also used to form cyclopropane and therefore the competing pathway results in a lower yield of propene *via* carbene insertion. In summary, propene is formed from both radical recombination and carbene insertion with a ratio of  $1.5 \pm 0.4 : 1 \pm 0.25$ , and propene and cyclopropane were formed from carbene insertion at a ratio of  $2.3 \pm 0.2 : 1 \pm 0.3$ .

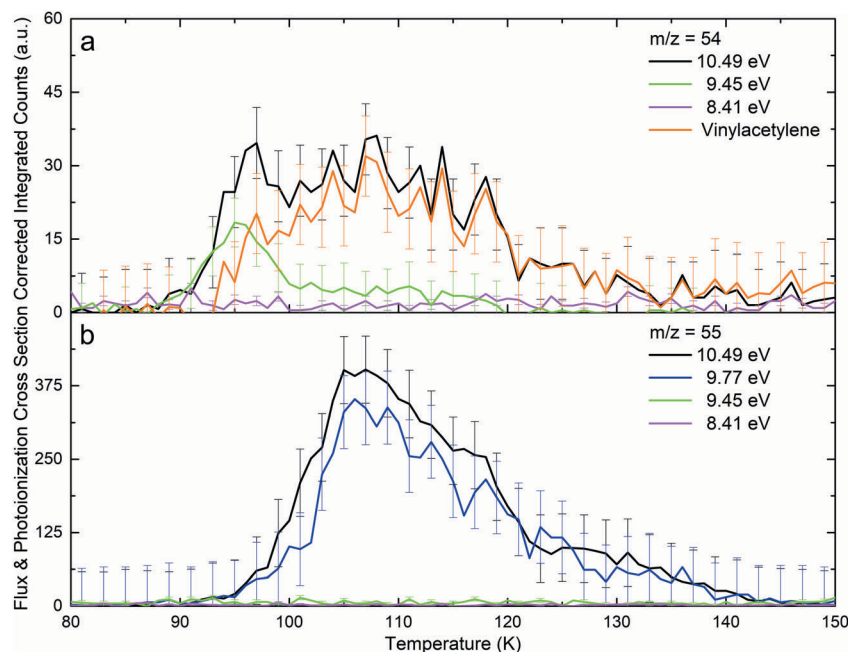
**4.1.3 C<sub>4</sub>H<sub>4</sub>.** Next, the vinylacetylene molecule's formation route was investigated taking into consideration the following pathways: reaction of dicarbon (C<sub>2</sub>) with ethylene (reaction (12)), or recombination of an ethynyl radical (C<sub>2</sub>H) with a vinyl radical (C<sub>2</sub>H<sub>3</sub>, reaction (13)). The reaction mechanism was probed *via* the mixed ices of D<sub>2</sub>-acetylene and <sup>13</sup>C<sub>2</sub>-ethylene, with a ratio of 1:1 (Fig. 7c). Detection of  $m/z = 54$  would confirm that vinylacetylene was synthesized from the reaction of dicarbon with

ethylene, and the detection of  $m/z = 55$  from this experiment reveals that vinylacetylene is produced *via* radical recombination of the ethynyl radical with the vinyl radical.

Fig. 10 shows that both  $m/z = 54$  and 55 were detected at a PI energy of 10.49 eV, but several tunable experiments were also needed to decipher which C<sub>4</sub>H<sub>4</sub> isomers this signal might be due to. Since, vinylacetylene has an IE of  $9.58 \pm 0.02$  eV any photoionization energy below this will no longer ionize this molecule, but all other C<sub>4</sub>H<sub>4</sub> species if they were produced. The detection of a signal for  $m/z = 54$  (PI = 9.45 eV) shows that it is not solely due to vinylacetylene, but with no signal detected at PI = 8.41 eV it was confirmed to be only due to 1,2,3-butatriene and vinylacetylene (Fig. 10a). Next, the non-detection of  $m/z = 55$  at PI = 9.45 eV shows that the radical recombination pathway can only synthesize the vinylacetylene isotopomer (Fig. 10b).

Again, the reactions to form the vinyl radical and ethynyl radical are endoergic and require 4.71 eV (454 kJ mol<sup>-1</sup>; reaction (7)) and 5.73 eV (553 kJ mol<sup>-1</sup>; reaction (3)), respectively. The ethynyl radical can lose another hydrogen atom forming dicarbon and this process is also endoergic by 4.90 eV (473 kJ mol<sup>-1</sup>; reaction (11)). The dicarbon molecule, which is a closed shell, has been shown to react barrierlessly with the ethylene molecule's double bond initially forming a reaction intermediate, which is then followed by isomerization to eventually form 1,2,3-butatriene or vinylacetylene, which agrees with the detection of both isomers.<sup>115</sup> The pathway forming vinylacetylene is exoergic by -6.09 eV (-588 kJ mol<sup>-1</sup>; reaction (12)).

Alternatively the recombination of the vinyl and ethynyl radicals to form vinylacetylene is also barrierless and exoergic by -5.90 eV (569 kJ mol<sup>-1</sup>; reaction (13)). Finally, the addition reaction of the vinyl radical with acetylene has been shown to



**Fig. 10** PI-ReTOF-MS flux and photoionization cross section corrected signals utilizing photoionization energies of 10.49 eV (black), 9.77 eV (blue), 9.45 eV (green), and 8.41 eV (pink) for (a)  $m/z = 54$  which was determined to be due to vinylacetylene (orange) and 1,2,3-butatriene (green) and (b)  $m/z = 55$ , which was due only to vinylacetylene, from the irradiated <sup>13</sup>C<sub>2</sub>H<sub>4</sub>-C<sub>2</sub>D<sub>2</sub> ice mixture.

have a barrier of 0.06 eV (6.2 kJ mol<sup>-1</sup>) and is exoergic by -0.17 eV (-16 kJ mol<sup>-1</sup>, reaction (14)).<sup>52</sup> Meanwhile, the addition of the ethynyl radical to ethylene does not have a barrier and is exoergic by -1.19 eV (-115 kJ mol<sup>-1</sup>, reaction (15)).<sup>52</sup> Here, the detection of  $m/z = 54$  and  $55$  in a ratio of  $1 \pm 0.3 : 11 \pm 3$  shows that radical recombination is strongly preferred for vinylacetylene synthesis (reaction (13)). Previous gas-phase experiments have shown that the reaction of dicarbon with ethylene preferentially formed 1,2,3-butatriene, which can then isomerize to vinylacetylene and may explain what is observed in the present experiment.<sup>115</sup> This agrees with the current results with the preferential formation of vinylacetylene in that the radical recombination pathway is heavily favored over insertion. In summary, vinylacetylene is synthesized through radical recombination and dicarbon reactions with a ratio of  $11 \pm 3 : 1 \pm 0.3$ .

## 5. Conclusions

Multiple studies have been conducted over the past decades to understand the astrochemical processes utilizing the C1 hydrocarbon methane (CH<sub>4</sub>) and the C2 hydrocarbons ethane (C<sub>2</sub>H<sub>6</sub>), ethylene (C<sub>2</sub>H<sub>4</sub>), and acetylene (C<sub>2</sub>H<sub>2</sub>). Although a thorough analysis of the different pathways to small hydrocarbons available through these C1 and C2 precursors has been attempted, an isomer specific study of the complex hydrocarbons that are formed, deep within these ices from interaction with GCRs, has been lacking until now. All of the hydrocarbon isomers detected here belong to a group in astrochemistry called complex organic molecules (COMs) – molecules having 6 or more atoms of carbon, hydrogen, nitrogen, and/or oxygen – which have been a key point of focus in recent years of laboratory astrophysics.<sup>2</sup> These COMs have been shown to be easily produced within the ice and not only through surface reactions.<sup>79</sup> The isomer specific identification of methylacetylene (CH<sub>3</sub>CCH), propene (CH<sub>3</sub>CHCH<sub>2</sub>), cyclopropane (c-C<sub>3</sub>H<sub>6</sub>), vinylacetylene (CH<sub>2</sub>CHCCH), 1-butyne (HCCCH<sub>2</sub>H<sub>3</sub>), 2-butyne (CH<sub>3</sub>CCCH<sub>3</sub>), 1,2-butadiene (H<sub>2</sub>CCCH(CH<sub>3</sub>)), and 1,3-butadiene (CH<sub>2</sub>CHCHCH<sub>2</sub>) provides insight into the complex hydrocarbon chemistry taking place in the ISM. Specifically, the investigation of gas-phase PAH formation has been a heavily studied area as PAHs are suggested to exist in large abundances in the ISM and account for much of the cosmic carbon,<sup>116–126</sup> and may have biological precursor relevance when nitrogen substituted.<sup>4,127,128</sup> Although some of the reaction pathways have been elucidated, several of the precursors' formation pathways needed for these reactions have remained elusive until now (Introduction).

For the C3 hydrocarbons detected here methylacetylene has been repeatedly shown to be a key precursor in PAH formation through bimolecular reactions with the phenyl radical (C<sub>6</sub>H<sub>5</sub>), in the gas phase, to produce indene – a bicyclic aromatic hydrocarbon.<sup>1,4–7</sup> Furthermore, the indene molecule has been shown to produce even more complex PAHs as large as chrysene (C<sub>18</sub>H<sub>12</sub>).<sup>129</sup> Also, pure methylacetylene has been shown to react with oxygen atoms to produce the complex oxygen-containing molecules such as C<sub>3</sub>H<sub>4</sub>O and C<sub>3</sub>H<sub>4</sub>O<sub>2</sub>, which consist of several

interesting isomers including some which have already been detected in the ISM.<sup>130</sup> It should be noted that allene (CH<sub>2</sub>CCH<sub>2</sub>), a second C<sub>3</sub>H<sub>4</sub> isomer that is nearly isoenergetic to methylacetylene, was not detected here, but is also very important for the formation of PAHs such as indene, 5-methyl-1*H*-indene, and 6-methyl-1*H*-indene.<sup>1,5,7,119</sup> While methylacetylene was the only isomer produced in the solid phase from methane ices bimolecular reactions under single collisions have determined that the reaction of the methylidyne radical (CH; X<sup>2</sup>Π) with ethylene forms only the allene isomer.<sup>131</sup> Therefore, the methylacetylene and allene isomers can be used to trace if the chemical evolution of different ISM environments have contributions from solid- or gas-phase chemistry, respectively. Additionally, the propene isomer has also been shown to produce RSFRs which can then yield PAHs, and also react with the phenyl radical producing indene (C<sub>9</sub>H<sub>8</sub>).<sup>1,12–14</sup> Propene was also recently shown to be a precursor in the formation of propylene oxide (c-C<sub>3</sub>H<sub>6</sub>O), which was the first chiral molecule detected in the ISM.<sup>132–134</sup> Also, the cyclopropane molecule, the only other possible C<sub>3</sub>H<sub>6</sub> isomer of propene, detected in these experiments represents the simplest cycloalkane. While cyclopropane is not a PAH precursor, it does prove that the complex hydrocarbons produced from solid methane ice can be cyclic.

Furthermore, the C4 hydrocarbon isomers selectively observed in the present manuscript have also been shown to produce PAHs. Vinylacetylene can react with the phenyl radical producing the prototypical PAH known as naphthalene (C<sub>10</sub>H<sub>8</sub>),<sup>18,135</sup> as well as with the *para*-tolyl radical (C<sub>6</sub>H<sub>4</sub>CH<sub>3</sub>) forming 2-methylnaphthalene (C<sub>11</sub>H<sub>10</sub>),<sup>19</sup> and finally with the *meta*-tolyl radical (C<sub>6</sub>H<sub>4</sub>CH<sub>3</sub>) creating 1-methylnaphthalene (C<sub>11</sub>H<sub>10</sub>).<sup>20</sup> Similarly, 1,3-butadiene forms multiple PAH precursors as well as PAH molecules. Intriguingly, the reaction of 1,3-butadiene with the ethynyl radical produces benzene, and the injection of 1,3-butadiene from dust grains greatly improves model abundances of benzene again showing a link between the solid and the gas phase.<sup>23</sup> Meanwhile, 1,3-butadiene's reaction with the phenyl radical generates 1,4-dihydronaphthalene (C<sub>10</sub>H<sub>10</sub>),<sup>41,136</sup> the reaction with the tolyl radical forms 5- and 6-methyl-1,4-dihydronaphthalene (C<sub>11</sub>H<sub>12</sub>),<sup>42</sup> and its reaction with the 1-naphthyl radical synthesizes dihydrophenanthrene (C<sub>14</sub>H<sub>12</sub>).<sup>137</sup> Furthermore, 1,3-butadiene can also react with the cyano radical to form pyridine (C<sub>5</sub>H<sub>5</sub>N),<sup>3,138</sup> and its reaction with pyridyl radicals (C<sub>5</sub>H<sub>4</sub>N) can produce 1,4-dihydro (iso)quinolone (C<sub>9</sub>H<sub>9</sub>N).<sup>3</sup> The other C<sub>4</sub>H<sub>6</sub> isomers detected here, 1,2-butadiene, 1-butyne, and 2-butyne, have also been shown to react with dicarbon to make RSFRs.<sup>139</sup>

The present manuscript shows that the coupling of traditional astrochemistry techniques with new highly sensitive analytical techniques can elucidate the complex hydrocarbon chemistry available in simple ISM ices. Fig. 11 depicts a global reaction scheme for the chemistry stemming from a methane ice. The formation routes of propane, ethane, ethylene, and acetylene, from methane ices, have been experimentally determined.<sup>140</sup> Now, the C3 hydrocarbons are complete with the addition of the C<sub>3</sub>H<sub>4</sub> and C<sub>3</sub>H<sub>6</sub> reaction pathways. Finally, the C4 hydrocarbons have been expanded upon as well with the addition of



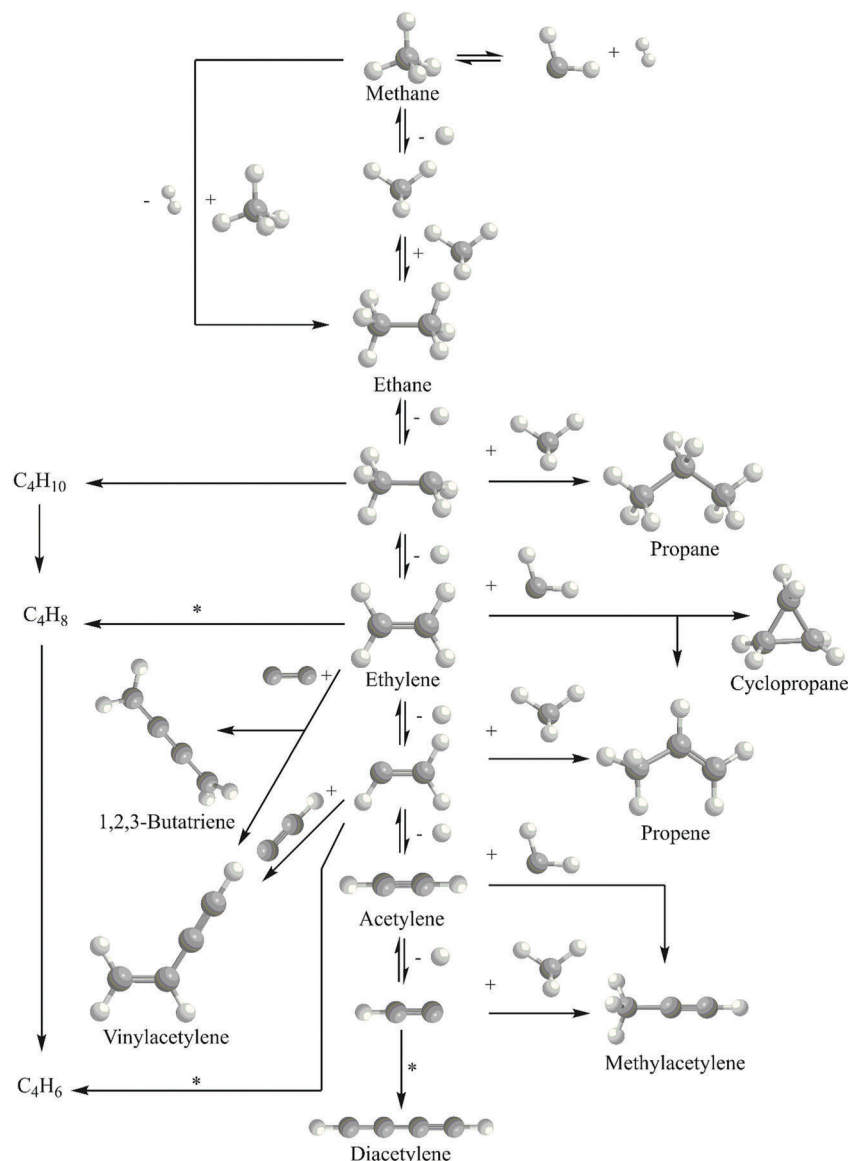


Fig. 11 The experimentally determined dominating reaction pathways for the formation of C2 hydrocarbons, as well as C<sub>3</sub>H<sub>4</sub>, C<sub>3</sub>H<sub>6</sub>, and C<sub>4</sub>H<sub>4</sub> isomers from methane ices (\* designates reaction pathways to be elucidated).

C<sub>4</sub>H<sub>4</sub> formation pathways. Furthermore, by studying the effects of adding other constituents to the ice, most importantly water, will help unravel the chemical complexity of hydrocarbon-containing ISM ices, and the current manuscripts provide the necessary first step in understanding these more realistic and complex mixtures. Finally, the results from this manuscript show a direct link between the solid phase, beyond surface reactions, and gas-phase chemistry taking place in the ISM. The inclusion of these solid-phase chemical reactions occurring within the ice rather than only on the surface has greatly enhanced the agreement of astrochemical models with observations for several COMs detected in the ISM.<sup>79</sup>

## Conflicts of interest

There are no conflicts to declare.

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