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A vacuum ultraviolet photoionization study on the formation of methanimine (CH₂NH) and ethylenediamine (NH₂CH₂CH₂NH₂) in low temperature interstellar model ices exposed to ionizing radiation†

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Methylamine (CH₃NH₂) and methanimine (CH₂NH) represent essential building blocks in the formation of amino acids in interstellar and cometary ices. In our study, by exploiting isomer selective detection of the reaction products *via* photoionization coupled with reflectron time of flight mass spectrometry (Re-TOF-MS), we elucidate the formation of methanimine and ethylenediamine (NH₂CH₂CH₂NH₂) in methylamine ices exposed to energetic electrons as a proxy for secondary electrons generated by energetic cosmic rays penetrating interstellar and cometary ices. Interestingly, the two products methanimine and ethylenediamine are isoelectronic to formaldehyde (H₂CO) and ethylene glycol (HOCH₂CH₂OH), respectively. Their formation has been confirmed in interstellar ice analogs consisting of methanol (CH₃OH) which is isoelectronic to methylamine. Both oxygen-bearing species formed in methanol have been detected in the interstellar medium (ISM), while for methanimine and ethylenediamine only methanimine has been identified so far. In comparison with the methanol ice products and our experimental findings, we predict that ethylenediamine should be detectable in these astronomical sources, where methylamine and methanimine are present.

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1. Introduction

During the last few decades, the methanimine molecule (CH₂NH) has received considerable attention from the astrochemistry and the astronomy communities due to its role as a potential precursor in the abiotic formation of amino acids such as glycine (NH₂CH₂COOH) – the simplest amino acid^{1,2} – in the interstellar medium (ISM). Although more than 80 amino acids were identified in carbonaceous chondrites like in the Murchison meteorite³ and recently glycine in the comet 81P/Wild 2,⁴ the understanding of their fundamental formation mechanisms is still in its infancy.^{5–10} Glycine, alanine (CH₃CH(NH₂)COOH), valine ((CH₃)₂CHCH(NH₂)COOH), proline (c-(NHCH₂CH₂CH₂CH)COOH), serine (HOCH₂CH(NH₂)COOH), and aspartic acid (HOC(O)CH₂-CH(NH₂)COOH) were identified *via* chromatography in the

room temperature residues of irradiated interstellar ice analogues containing water (H₂O), ammonia (NH₃), methanol (CH₃OH), hydrogen cyanide (HCN), carbon monoxide (CO), and carbon dioxide (CO₂).^{10–13} These results elucidated a radiation induced low temperature formation pathway toward glycine commencing with the decomposition of methylamine (CH₃NH₂) *via* production of CH₂NH₂ together with suprathreshold hydrogen atoms. The latter react with CO₂ to form the hydroxycarbonyl radicals (HOCO), which then recombines with the aminomethyl radicals (CH₂NH₂) to form glycine. This pathway was also supported by quantum chemistry calculations.¹⁴ Alternatively, Kaiser *et al.*¹⁰ discussed a formation pathway toward glycine (NH₂CH₂COOH) *via* NH₃ and acetic acid (CH₃COOH) – the latter was previously detected in irradiated CO₂ and methane (CH₄) ices^{15,16} – *via* radical – radical recombination of radiolytically generated amino (NH₂) and hydroxycarbonylmethyl radicals (CH₂COOH).

The CH₂NH molecule itself – also called formaldimine due to the isoelectronic structure with formaldehyde (H₂CO) – was first detected in the hot core Sagittarius B2 toward the galactic center through the 64 m radio telescope *via* the 1₁₀–1₁₁ multiplet¹⁷ with a column density of 3 × 10¹⁴ cm⁻². Follow-up searches consolidate this finding^{18–22} and confirmed the presence of CH₂NH in L183,²³ Orion-KL,^{1,24} W51, Orion 3N, G34.3+0.15,¹

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and G19.61-0.23,²⁵ as well as in the circumstellar shell of IRC+10216.²⁶ Methanimine was also detected in the ultraluminous infrared galaxy Arco ARP 220.²⁷ In the solar system, the atmosphere of Saturn's moon Titan also reveals an abundance of CH₂NH as discovered by the Cassini T5 flyby.²⁸ Furthermore, methylamine has been detected in dust samples collected by the Stardust mission⁴ and was recently also observed on the comet 67P/Churyumov-Gerasimenko.²⁹ In summary, we can find CH₂NH in every environment in the interstellar medium which underlines its importance in the interstellar chemistry.

However, despite the potential importance of CH₂NH, the underlying formation pathways are still not resolved. In the gas phase, bimolecular neutral-neutral reactions of methyldiyne (CH) with NH₃ have been proposed to synthesize CH₂NH along with atomic hydrogen (H).²⁶ In laboratory experiments, Michael *et al.*³⁰ detected traces of CH₂NH through photolysis of CH₃NH₂ by non-monochromatic ultra violet (UV) radiation around 220 nm. However, formation rates through these pathways are too low and cannot explain the observed fractional abundances of CH₂NH in the interstellar medium.^{23,31} For instance, toward the cold cloud L183, fractional abundances of CH₂NH were detected at a level of 8.1×10^{-10} ,^{1,23} but astrochemical models predict fractional abundances about one to two orders of magnitude below the actual observations. Therefore, gas phase chemistry alone cannot account for the formation of interstellar CH₂NH, and critical production routes are still lacking.

An alternative source for CH₂NH can be found in formation routes in condensed matters, *i.e.* within interstellar ices containing NH₃ and CH₄ exposed to ionizing radiation. Here, methanimine has been detected tentatively in broad band ultraviolet (UV) irradiated interstellar model ices carrying CH₄ and NH₃.^{32,33} Photolysis of CH₃NH₂ ice has also been studied by Bossa *et al.*³⁴ tentatively assigning CH₂NH *via* its symmetric C–H stretching vibration at 3144 cm⁻¹ appearing as a shoulder of the hydrogen-bonding mode of the CH₃NH₂ parent. Woon *et al.*¹⁴ discussed the formation of CH₂NH along with CH₃NH₂ *via* a sequential hydrogenation of HCN, which was confirmed experimentally by Theule *et al.*³⁵ Kim *et al.*³⁶ demonstrated that a radical-radical recombination between methyl (CH₃) and amide (NH₂) in CH₄-NH₃ bearing ices followed by radiolysis leads ultimately to HCN *via* CH₃NH₂ and CH₂NH intermediates.

The aforementioned considerations revealed that an experimental elucidation of the synthetic pathways to CH₂NH in low temperature interstellar analog ices exposed to ionizing radiation is still in its infancy. Nevertheless, they provide a significant potential to account for the missing CH₂NH source in the ISM. Previously analytical techniques have been used to detect CH₂NH *via* low temperature fourier transform infrared spectroscopy (FTIR) of the exposed ices.³⁴ Additionally, quadrupole mass spectrometry (QMS) coupled with electron impact ionization has been used in that study to probe CH₂NH subliming in the post irradiation phase into the gas phase. These approaches hold significant complications. FTIR spectroscopy represents an ideal tool to investigate the processing and decay kinetics of 'small' molecules such as CO, H₂O, CH₃OH, CO₂, CH₄ and

H₂CO, along with NH₃ – these molecules have been detected on interstellar grains. However, the ability of infrared spectroscopy to provide useful information for the detection of complex organics formed within the ices is quite limited. Infrared spectroscopy allows the identification of functional groups of organics. This information does not always identify individual molecules since the functional groups of, for instance, amines (R–NH₂) and imines (RHC=NH) portray similar group frequencies in the range of 1650 cm⁻¹ to 1500 cm⁻¹ and 3500 cm⁻¹ to 3300 cm⁻¹.³⁷ Therefore, the exclusive assignment of a newly formed molecule based on infrared bands in an unknown mixture of organics is rarely scientifically sound. Using QMS coupled with electron impact ionization to ionize the subliming molecules of the irradiated ices during the temperature programmed desorption (TPD) exploits often an electron impact ionizer operating at 70 to 100 eV electron energy. This energy range does not only ionize molecules, but also results in a significant fragmentation of the parent ion. This might even result in the absence of the molecular parent ion. Furthermore, the fragment ions of structural isomers often overlap making it difficult to decipher or even to discriminate between structural isomers. The exploitation of soft ionization with low energy electrons of a few electron volt kinetic energy has a few advantages. However, a voltage drop across the filaments of typically 1.0 eV results in electrons with a relatively broad energy distribution thus making it difficult to selectively ionize structural isomers. Therefore, alternative analytical techniques are required to elucidate the formation of CH₂NH in low temperature interstellar ice analogues.

In this work, we use single photon ionization (PI) coupled with a reflection time-of-flight mass spectrometer (ReTOF-MS) to detect and to distinguish structural isomers of organic molecules subliming into the gas phase during temperature programmed desorption (TPD) after exposing the ices to ionizing radiation. In comparison to traditional electron impact ionization, PI-ReTOF-MS – utilizing photon energies close to the ionization threshold of the molecules – forms gas phase ions without significant internal energy and hence ideally without fragmentation.^{38,39} Likewise, employing distinct photon energies of 9.93 eV, 9.10 eV, and 8.17 eV as conducted in the present work, this technique can be utilized to selectively discriminate between the structural isomers of organic molecules as previously demonstrated.⁴⁰ The present study demonstrates for the first time unambiguously that CH₂NH can be synthesized in low temperature (5 K) interstellar model ices of CH₃NH₂, when exposed to ionizing radiation such as energetic electrons. These energetic electrons simulate secondary electrons released during the passage of galactic cosmic rays through interstellar ices.⁴¹ The present laboratory simulation experiments mirror the exposure of the interstellar ices in molecular clouds to galactic cosmic rays during a life time of a molecular cloud of typically 10⁵ years. Considering that molecular clouds constitute the nurseries of stars and planetary systems,^{42–44} the identification of CH₂NH along with ethylenediamine (NH₂CH₂CH₂NH₂) in our experiments suggests that these molecules may have been at least partially incorporated into our solar system from interstellar matter *via* circumstellar disks. Finally, considering the

isoelectronicity between CH_3NH_2 and CH_3OH , the formation pathways of the CH_2NH and $\text{NH}_2\text{CH}_2\text{CH}_2\text{NH}_2$ molecules are compared to those of the H_2CO and ethylene glycol ($\text{OHCH}_2\text{CH}_2\text{OH}$) in interstellar model ices exposed to ionizing radiation.^{41,45–47}

2. Experimental methods

The experiments were performed at the W. M. Keck Research Laboratory in Astrochemistry. The experimental setup consists of a contamination-free stainless steel ultra-high vacuum chamber (UHV) operating at a base pressure of a few 10^{-11} Torr backed by turbo molecular pumps and dry oil-free scroll pumps. A polished polycrystalline silver mirror within the chamber is coupled to a cold finger cooled to 5.5 ± 0.1 K using a UHV compatible closed-cycle helium compressor (Sumitomo Heavy Industries, RDK-415E). The target can be moved in two degrees of freedom, translation in the vertical axis and rotation in the horizontal plane. Methylamine gas (Sigma Aldrich; 99%) was deposited onto the cooled silver wafer *via* a glass capillary at a base pressure of 5.5×10^{-9} Torr for 7 min until the desired ice thickness was achieved. This ice growth was monitored online and *in situ* *via* laser interferometry measurements. In this method, the HeNe laser ($\lambda = 632.8$ nm; CVI Melles-Griot; 25-LHP-230) reflections of the silver mirror and the ice surface introduce an interference pattern. The beam was reflected into a photodiode and the pattern is measured using a picoammeter (Keithley 6485). The recorded fringes result in the thickness of the ice *via* eqn (1), which describes the thickness (d) of an ice in dependence of the number of interference fringes (m), the laser wavelength (λ), and the angle of incidence of the laser beam ($\theta = 4^\circ$):^{48–50}

$$d = \frac{m\lambda}{2\sqrt{n^2 - \sin^2\theta}} \quad (1)$$

A refractive index (n) for CH_3NH_2 ice of 1.40 ± 0.30 is derived from the density ($\rho = 0.785$ g cm^{-3}) and refractive index ($n_f = 1.371$) of liquid CH_3NH_2 *via* the Lorentz–Lorentz relation^{51,52} with the Lorenz coefficient (L) to be constant over a fixed wavelength:

$$L\rho = \frac{n_f^2 - 1}{n_f^2 + 2} \quad (2)$$

With the density of CH_3NH_2 ice of $\rho = 0.85$ g cm^{-3} ,^{53,54} the thickness of the prepared CH_3NH_2 ice is 200 ± 50 nm. After the deposition with a rate of 30 ± 5 nm min^{-1} , an area of 1.0 ± 0.1 cm^2 of CH_3NH_2 ice was then irradiated by an electron gun (specs, PU-EQ22) with 5 keV electrons for 15 min with an electron current of 20 nA at an angle of incidence of 70° relative to the surface normal. Galactic cosmic rays (GCR) consist mainly of protons with kinetic energy up to PeV. Interaction with interstellar ices and ionization of molecules lead to an energy loss of the GCRs and result in secondary electrons which subsequently ionize further molecules generating electron cascades with energies in the range of eV–keV.⁵⁵ Therefore, the 5 keV electrons mimic efficiently the irradiation by GCRs. By utilizing the CASINO 2.42 software,⁵⁶ an average penetration depth of 180 ± 80 nm is achieved (Table S1, ESI[†]). The average

dose absorbed is determined to be 1.0 ± 0.1 eV molecule⁻¹. During the irradiation, the chemical evolution of the CH_3NH_2 ice was monitored online and *in situ* using FTIR (Nicolet 6700, MCT-A) in the range of 4500 cm^{-1} – 500 cm^{-1} , with a resolution of 4 cm^{-1} , at a reflection angle of 45° for absorption–reflection–absorption mode. Once the irradiation was completed, the ice was held at 5.5 K for additional 30 min before starting temperature programmed desorption (TPD) by heating the substrate from 5.5 K to 320 K at a rate of 0.5 K min^{-1} .

To detect the subliming molecules with the PI-ReTOF-MS, the neutral molecules are subliming during TPD and are ionized by pulsed vacuum ultraviolet (VUV) laser light close to 2 mm in front of the ice sample. The ions produced are mass-resolved and detected by a multichannel plate in a dual chevron configuration and then amplified by a fast preamplifier (Ortec 9305) and shaped with a 100 MHz discriminator. Finally, the spectra are recorded using a computer-controlled multichannel scaler (FAST ComTec, P7888-1E) with 4 ns bin width triggered at 30 Hz using a pulse delay generator (Quantum Composer, 9518) and 3600 sweeps per mass spectrum per 1 K increase in temperature during TPD.

VUV radiation is generated by difference-frequency mixing ($2\omega_1 - \omega_2$) in a noble gas (krypton (Kr) or xenon (Xe)) using two dye lasers (Sirah Lasertechnik, Model Cobra-Stretch, and Precision Scan) each pumped with a Nd:YAG laser (Spectra-Physics, Models PRO-270-30). The laser wavelengths were measured using a WaveMaster laser wavelength meter (COHERENT). In order to record and monitor VUV light, a detection system consisting of a Faraday cup and a diode calibrated by the national institute of standards and technology (NIST) is placed behind the detection region. This tunable VUV photon source allows soft-ionization of molecules with almost no fragmentation.⁴⁰ In this work, we used a mixture of Rhodamine 610 and Rhodamine 640 in the first dye laser, pumped with 532 nm, to generate 606.948 nm, which is frequency tripled to $\omega_1 = 202.316$ nm for the two-photon resonance of krypton (Kr). Adding the second laser at a wavelength of $\omega_2 = 532$ nm results in a VUV energy of 9.93 eV (124.9 nm). For the second VUV energy of 9.10 eV, the first dye laser with Coumarin 450 as dye is pumped with 355 nm to achieve 445.132 nm which result in $\omega_1 = 222.566$ nm by frequency doubling for the two-photon resonance in xenon (Xe). The second dye laser with Rhodamine 610/Rhodamine 640 is pumped with 532 nm to generate 607 nm. The final photoionization energy used is 8.17 eV (151.8 nm). It is generated by different frequency mixing of $\omega_1 = 249.628$ nm and $\omega_2 = 703$ nm in the xenon gas (Xe). The first dye laser is prepared with Coumarin 503 and pumped with 355 nm to achieve a wavelength of 499.265 nm before frequency doubling. The second wavelength is generated by the laser dye LDS 867 pumped with 532 nm. An overview of the photon ionization energies and their preparation can be found in Table 1. Calibration measurements for the ReTOF reveal a potential lowering of the ionization energies of the subliming molecules by up to 0.03 to 0.04 eV due to the static electric field induced Stark effect.⁵⁷ This has been confirmed by separate calibration experiments using molecules with known ionization potentials, such as methyl bromide (CH_3Br) (IE = 10.540 ± 0.003 eV⁵⁸), dibromochloromethane (CHBr_2Cl)

Table 1 Parameter for the vacuum ultraviolet light generation used in the present experiments. The uncertainty for VUV photon energies is 0.01 eV

	Photoionization energy (eV)	9.93	9.10	8.17
	Flux (10^{11} photons s^{-1})	10 ± 1	6 ± 1	15 ± 1
$(2\omega_1 - \omega_2)$	Wavelength (nm)	124.9	136.2	151.8
ω_1	Wavelength (nm)	202.316	222.566	249.628
Nd:YAG (ω_1)	Wavelength (nm)	532	355	355
Dye laser (ω_1)	Wavelength (nm) – dye laser	606.948	445.132	499.265
Dye		Rh 610/Rh 640	C450	C503
ω_2	Wavelength (nm)	532	607	703
Nd:YAG (ω_1)	Wavelength (nm) – Nd:YAG	532	532	532
Dye laser (ω_1)	Wavelength (nm) – dye laser	—	607	703
Dye		—	Rh 610/Rh 640	LDS 867
	Nonlinear medium	Kr	Xe	Xe

(IE = 10.59 ± 0.01 eV⁵⁹) and hexafluoropropylene (C_3F_6) (IE = 10.60 ± 0.03 eV⁶⁰).

3. Computational methods

The hybrid density functional B3LYP^{61–64} with the cc-pVTZ basis set was employed to optimize geometries and harmonic frequencies of the species and their cations. Afterwards, their coupled cluster^{65–68} CCSD(T)/cc-pVDZ, CCSD(T)/cc-pVTZ, and CCSD(T)/cc-pVQZ energies were calculated and extrapolated to complete basis set limits,⁶⁹ CCSD(T)/CBS, with B3LYP/cc-pVTZ zero-point energy corrections. The energy difference between the ionic and neutral states with similar geometries gives the adiabatic ionization energy. The GAUSSIAN09 program⁷⁰ was utilized in the electronic structure calculations. Table 2 presents the calculated adiabatic ionization energies for two

sets of isomers $C_2N_2H_8$ along with the relative energies of the structural isomers considered in this study and ionization energy values from the literature. A comparison between theoretical and experimental results as a benchmark calculation reveals that the calculations overestimate the ionization energies by 0.03 eV to 0.05 eV.^{71–73} Molecular geometries as well as vibrational modes for the $C_2N_2H_8$ isomers are presented in the supplement material (Tables S2 and S3, ESI†).

4. Results

Fig. 1 presents the FTIR spectra of CH_3NH_2 before (black line) and after (red line) the irradiation for the high ($3500\text{--}2700\text{ cm}^{-1}$) and low ($1700\text{--}800\text{ cm}^{-1}$) energy regions. All absorptions can be attributed to CH_3NH_2 as presented by previous studies.^{34,74} Their assignments are compiled in Table 3. The region from 3400 to

Table 2 Calculated adiabatic ionization energies (IE) and the relative energies (E_{rel}) of distinct $C_2N_2H_8$ and $C_2N_2H_8$ isomers. The '3' superscript indicates a triplet state

Structure	Molecular formula	Species	IE ^a (eV)	E_{rel} ^b (eV)	Literature value IE, (eV)
	CH_2NH	Methanimine	9.94	0.00	9.88 ± 0.07^c
	3CH_3N	Methylimidogen	12.00	2.34	
	$CHNH_2$	Aminomethylene	8.22	1.54	
	3CHNH_2	Aminomethylene	10.94	3.01	
	$NH_2CH_2NHCH_3$	<i>N</i> -Methyl-methanediamine	8.23	0.23	
	$NH_2CH_2CH_2NH_2$	Ethylenediamine	8.27	0.00	8.42 ± 0.04^d
	$CH_3NHNHCH_3$	1,2-Dimethylhydrazine	7.18	1.28	7.8 ± 0.1^e
	$(CH_3)_2NNH_2$	1,1-Dimethylhydrazine	7.24	1.16	7.29 ± 0.05^f

^a Relative ionization potential by CCSD(T)/CBS with B3LYP/cc-pVTZ zero-point energy correction in eV. ^b Relative energy by CCSD(T)/CBS with B3LYP/cc-pVTZ zero-point energy correction in eV. ^c Taken from Tarasenko *et al.*, 1986.⁹⁶ ^d Taken from Wei *et al.*, 2006.⁸⁰ ^e Taken from Dibeler *et al.*, 1959.⁹⁷ ^f Taken from Moet-Ner *et al.*, 1984.⁹⁸

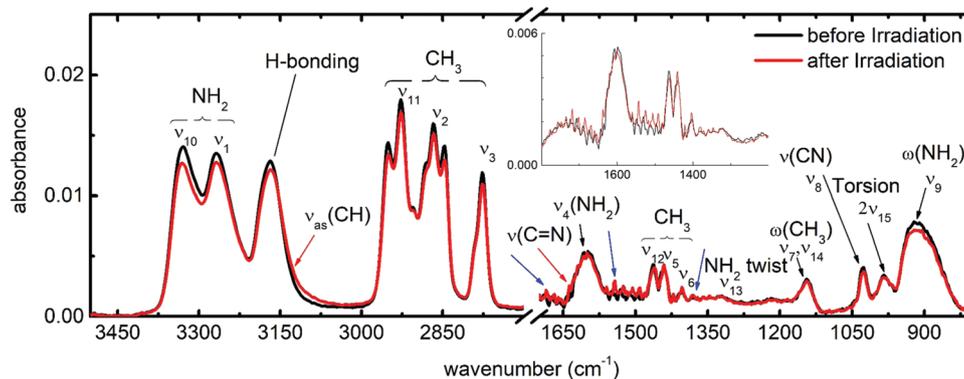


Fig. 1 FTIR spectra before (black) and after (red) irradiation of the CH_3NH_2 ice for the high and low energy regions. Besides a loss in the total absorbance after irradiation only a few new features are observed. The shoulders around 3140 cm^{-1} and 1635 cm^{-1} (red arrows) could be tentatively attributed to CH_2NH . Further features at 1380 cm^{-1} , 1540 cm^{-1} , and 1680 cm^{-1} (blue arrows) cannot be attributed to one particular molecule. Since no features can be found between 2700 and 1700 cm^{-1} , these data are not presented, for better display in the relevant regions. The inset expands the low-intensity region of the spectra.

3100 cm^{-1} is governed by the NH_2 antisymmetric (ν_1) and symmetric (ν_{10}) stretching vibrations, while the region of $3000\text{--}2700\text{ cm}^{-1}$ connects to the two degenerate (ν_{11} , ν_2) and the symmetrical (ν_3) CH_3 stretching vibrations of CH_3NH_2 . The multiple peak structures and peak shift of these modes are characteristic for Fermi resonances. The feature at 3170 cm^{-1} is interpreted by Durig *et al.*⁷⁴ as hydrogen bonding to the nitrogen electron lone pair. At lower energies, features for the scissor mode (ν_4) of NH_2 and the degenerate (ν_{12} , ν_5) and symmetric (ν_6) CH_3 deformations are visible followed by the very weak NH_2 twisting (ν_{13}). From 1250 cm^{-1} , the C–N (ν_8) stretching as well as the first overtone of the torsion (ν_{15}) can be found surrounded by the NH_2 wagging (ν_9) and CH_3 rocking (ν_7 , ν_{14}) modes. The modes found in this study agree with the findings of Durig *et al.*⁹⁶ as presented in Table 3.

For the irradiated sample, only minor differences can be detected considering the low irradiation dose of only $1.0 \pm 0.2\text{ eV molecule}^{-1}$. The intensities of the CH_3NH_2 fundamental vibrations are reduced compared to the unirradiated sample by $8 \pm 1\%$ (average decrease of each fundamental band area), revealing that $0.09 \pm 0.01\text{ CH}_3\text{NH}_2$ molecules eV^{-1} are reacting during the radiation exposure. In the region of $3000\text{--}3200\text{ cm}^{-1}$, a shoulder at 3140 cm^{-1} becomes visible. This shoulder can be attributed tentatively to the antisymmetric C–H (ν_{as}) stretching mode of CH_2NH .³⁵ A second shoulder can be found at 1635 cm^{-1} in the NH_2 scissor mode (ν_4). This feature might be attributed to the C=N double bond stretching of CH_2NH (Table 3). Further, new modes around 1650 cm^{-1} , 1550 cm^{-1} , and 1380 cm^{-1} (blue arrows) could be attributed to several possible functional groups. Around 1650 cm^{-1} , mostly C=C and C=N stretching vibrations can be found, while features around 1550 cm^{-1} are most likely associated with the amine functional group *via* NH_2 scissor or NH deformation. The features around 1380 cm^{-1} are typical for alkane functional groups visible by CH deformation modes. The newly formed molecules consisting of C, N, and H can be attributed to these features. However, due to the weak appearance and overlaps of functional groups for different molecules, it is difficult to identify individual molecules. Taking into account the energy loss of the electrons while penetrating the ices (Table S1, ESI[†]), $(25 \pm 3) \times 10^{15}$ CH_3NH_2 molecules are destroyed

Table 3 Infrared absorption features of CH_3NH_2 ⁷⁴ and CH_2NH ³⁴

Wavenumber (cm^{-1}) from the literature	Wavenumber (cm^{-1}) this study	Vibration mode	Assignment
CH_3NH_2			
3332	3331	NH_2 antisymmetric stretching	ν_{10}
3260	3260	NH_2 symmetric stretching	ν_1
3191	3170	H bonding	
2942	2945	CH_3 degenerate stretching	ν_{11}
2881	2875	CH_3 degenerate stretching	ν_2
2793	2785	CH_3 symmetric stretching	ν_3
1636	1606	NH_2 scissoring	ν_4
1500	1493	CH_3 degenerate deformation	ν_{12}
1467	1465	CH_3 degenerate deformation	ν_5
1441	1445	CH_3 symmetric deformation	ν_6
1353	1356	NH_2 twisting	ν_{13}
1182	1142	CH_3 rocking	ν_7 , ν_{14}
1048	1030	CN stretching	ν_8
1005	983	Torsion first overtone	$2\nu_{15}$
955	917	NH_2 wagging	ν_9
CH_2NH			
3144	3120	CH antisymmetric stretching	ν_{a}
1662	1643	C=N stretching	$\nu(\text{C}=\text{N})$

during irradiation in an area of 1 cm^2 . If each destroyed molecule is converted to CH_2NH , 0.09 ± 0.01 molecules of CH_2NH are formed per eV. Naturally, this value can only be considered as an upper limit for the CH_2NH formation, since other molecules are formed as well based on the observation of the aforementioned functional groups and the Re-TOF-MS data as discussed below. In conclusion, the FTIR results show the preparation and processing of CH_3NH_2 ice. However only tentatively results can be found for the products formed.

Compared to FTIR, the PI-ReTOF-MS detection method is significantly more sensitive and can discriminate between structural isomers – information which hardly can be extracted by alternative techniques in astrophysical ice simulation experiments.^{16,48,75–77} The PI-ReTOF-MS data are compiled in Fig. 2 for distinct ionization energies of 9.93 eV, 9.10 eV, and 8.17 eV while the corresponding TPD graphs for selected m/z ratios critical to the present study are presented in Fig. 3. Fig. 2a and b represent the PI-ReTOF-MS data of the unirradiated (blank) and irradiated ices, respectively, recorded with the highest photon energy of 9.93 eV. The most intense signals in the control experiment and the irradiated ice sample appear at mass-to-charge ratios (m/z) of 30, 31, 32, and 33 in the temperature range from 100 K to 120 K. The most intense ion counts as presented in Fig. 3a can be attributed to the parent molecule CH_3NH_2 (CH_3NH_2 ; 31 amu; IE = 8.97 ± 0.02 eV).^{78,79} Since CH_3NH_2 has an ionization energy of 8.97 eV, it cannot be observed at a photon energy of 8.17 eV (Fig. 2d). For photon energies above 9 eV, the appearance of the ion at $m/z = 30$ could be assigned to CNH_4^+ ($m/z = 30$), which results from atomic hydrogen loss of CH_3NH_2 (CH_3NH_2 ; 31 amu). Furthermore, the ion counts at $m/z = 32$ can be explained by protonated CH_3NH_2 (CH_3NH_3^+) and the naturally occurring ^{13}C -substituted CH_3NH_2 ($^{13}\text{CH}_3\text{NH}_2$); the protonated ^{13}C -substituted CH_3NH_2 ($^{13}\text{CH}_3\text{NH}_3^+$) can be linked to $m/z = 33$. A noticeable smaller feature can be observed in the pristine and irradiated systems at $m/z = 45$ desorbing at the same temperature as CH_3NH_2 . This signal can be attributed to ionized ethylamine ($\text{CH}_3\text{CH}_2\text{NH}_2$) and/or dimethylamine (CH_3NHCH_3). These molecules have ionization energies of 8.9 eV and 8.2 eV (Table 4), respectively, and can be attributed to tracer impurities in the

CH_3NH_2 gas (99%). Furthermore, the ion signal at $m/z = 59$ can be related to propylamine ($\text{CH}_3\text{CH}_2\text{CH}_2\text{NH}_2$), 2-propanamine ($(\text{CH}_3)_2\text{CHNH}_2$), and/or *n*-methyl ethanamine ($\text{CH}_3\text{CH}_2\text{NHCH}_3$) which have ionization energies of 8.78 eV, 8.72 eV, and 8.15 eV, respectively (Table 4).

After the irradiation, several new mass-to-charge ratios were observed, which were absent in the blank experiment. The TPD traces are presented in Fig. 3. Due to the simplicity of the ice, the resulting molecules after the irradiation can only be composed of three elements: hydrogen (H), nitrogen (N), and carbon (C). The new signal at $m/z = 29$ appearing between 106 K and 124 K can only be explained by four different isomers of CNH_3 : CH_2NH , CHNH_2 , triplet methylimidogen ($^3\text{CH}_3\text{N}$), and triplet aminomethylene ($^3\text{CHNH}_2$) (Table 2). The latter two isomers have ionization energies well above 9.93 eV and cannot be detected in this study. Therefore, they can be ruled out as source for the trace of $m/z = 29$. Based on the ionization energies of 9.88 ± 0.07 eV for CH_2NH and 8.22 ± 0.05 eV for CHNH_2 , the two isomers can be distinguished by tuning the photoionization energy between these two ionization energies. Fig. 3a presents the TPD spectrum of $m/z = 29$ at 9.93 eV and 9.10 eV photoionization energy. The strong clear peak at 120 K for 9.93 eV vanishes for 9.10 eV (Fig. 3a). Based on these findings, $m/z = 29$ can be assigned to CH_2NH . Therefore, we can conclude that CH_2NH is identified as the product in the radiolysis of CH_3NH_2 .

The remaining TPD traces in Fig. 3b reveal sublimation maxima at higher temperatures between 135 K and 225 K. A mass-to-charge ratio of 60 represents the heaviest ion detected.

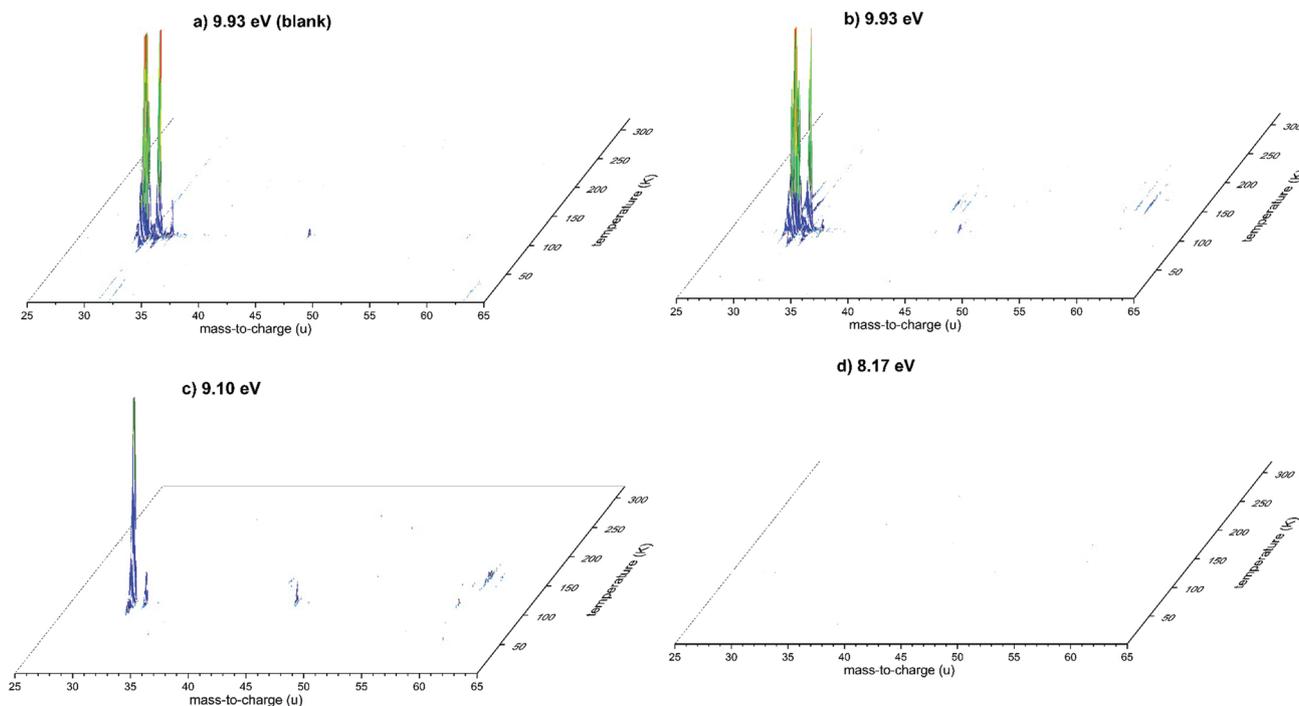


Fig. 2 PI-ReTOF-MS spectra of the sublimating molecules recorded for the irradiated CH_3NH_2 ices obtained at distinct ionization energies. Panel (a) shows the non-irradiated CH_3NH_2 ice (blank) while panel (b), (c) and (d) present the irradiated ice analyzed at 9.93 eV, 9.10 eV and 8.17 eV photoionization energy, respectively.

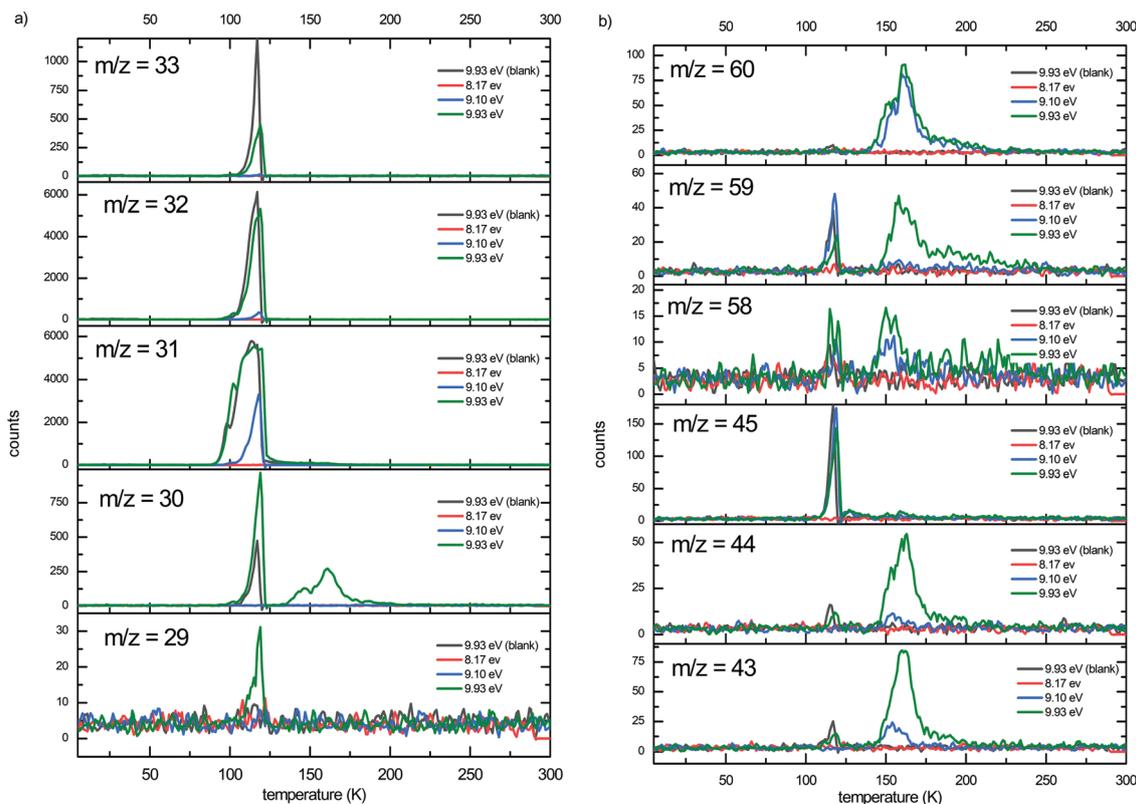


Fig. 3 TPD traces of lower (a) and higher mass (b) ranges. The m/z ratio and ionization energy of each trace are given in the legend.

It can be attributed to the molecular formula $C_2N_2H_8$ that can be assigned to four possible isomers: $NH_2CH_2CH_2NH_2$, N -methyl-methanediamine ($NH_2CH_2NHCH_3$), 1,1-dimethylhydrazine ($(CH_3)_2NNH_2$), and 1,2-dimethylhydrazine ($CH_3NHNHCH_3$). They can be distinguished *via* their ionization energies (Table 2). Ethylenediamine has the highest ionization energy of 8.27 eV (calculated) and 8.42 eV (experimental) followed by $NH_2CH_2NHCH_3$ with 8.23 eV. Taking the energy correction of the Stark shift (0.04 eV) and the offset of the theoretical ionization energy (0.04 eV) into account, the ionization energies of $NH_2CH_2NHCH_3$ and $NH_2CH_2CH_2NH_2$ are reduced to 8.15 eV and 8.20 eV, respectively. A photon energy of 8.17 eV is employed to distinguish between both isomers. As presented in Fig. 3b, no ion counts are detected at 8.17 eV, revealing that most likely $NH_2CH_2CH_2NH_2$ is formed during irradiation, which effectively contributes to ion counts at $m/z = 60$ in the 9.93 eV experiment. However, due to the error of ± 0.05 eV on the calculated ionization energies, we cannot rule out $NH_2CH_2NHCH_3$ entirely, yet. Nevertheless, we find ion masses at $m/z = 59, 58, 44, 43,$ and 30 which simultaneously subliming with $m/z = 60$. In comparison to an ethylenediamine photoionization study by Wei *et al.*,⁸⁰ we can relate the ion masses in this study to the photofragments of $NH_2CH_2CH_2NH_2$ (Table 4). These findings confirm the formation of $NH_2CH_2CH_2NH_2$.

In summary, the PI-ReTOF-MS study reveals two newly formed molecules in the CH_3NH_2 ices exposed to ionizing radiation: CH_2NH and $NH_2CH_2CH_2NH_2$. Integrating over the signals in the TPD traces of $m/z = 32$ with and without irradiation helps to extract the fraction of the non-radiolyzed CH_3NH_2

molecules to be $92 \pm 4\%$. This coincides nicely with the calculated loss of about $8 \pm 1\%$ from the infrared study.

The previous experiment from Bossa *et al.* already proposed the formation of CH_2NH . In their photolysis study, they found small features in their FTIR study around 1722 cm^{-1} and 1683 cm^{-1} which they postulated to originate from amides or aldehydes. Therefore, they related the $m/z = 59$ during sublimation using QMS to N -methyl formamide. In their experiment, they used UV flux of $10^{15}\text{ photons s}^{-1}\text{ cm}^{-2}$. This flux converts to about $14\text{ eV molecule}^{-1}$,⁸¹ which is in the same region than the energy dose used in this work. Their findings, however, only suggest the existence of N -methyl formamide, based on the QMS data. An isotope study or an analysis of the isomer structure of $m/z = 59$ as we present it in this study has not been done in their study.

5. Discussion

Having confirmed that CH_2NH and $NH_2CH_2CH_2NH_2$ are formed in the irradiated ices, we are discussing now the possible formation pathways. Holtom *et al.*¹³ revealed that CH_3NH_2 can undergo unimolecular decomposition initiated by energy transfer from the impinging electrons. In particular, two hydrogen loss channels can result in either methylamidogen (CH_3NH) (3a) or CH_2NH_2 (4a) (see also Fig. 4); in the present experiments. Each of these radicals can emit a hydrogen atom synthesizing CH_2NH (3b and 4b). These routes are highly endoergic with the

Table 4 Species observed and discussed in this work along with their molecular formula, mass-to-charge ratio (m/z), ionization energy (IE), appearance energy (AE) of the observed fragments, and the observed peak sublimation temperature, as well as the Gibbs free energy $\Delta_r G^\circ$

Species	Chemical formula	m/z	IE (eV)	Sublimation peak (K)	$\Delta_r G^\circ$ (kJ mol ⁻¹)
Hydrogen	H	1			216 ^d
Methanimine	CH ₂ NH	29	9.94 ^a	115	97 ^d
Methylimidogen	³ CH ₃ N	29	12.00 ^a		
Aminomethylene	CHNH ₂	29	8.22 ^a		
Aminomethylene	³ CHNH ₂	29	10.94 ^a		
Methylamidogen	CH ₃ NH	30			187 ^d
Aminomethyl	CH ₂ NH ₂	30			160 ^d
Methylamine	CH ₃ NH ₂	31	8.97 ^b	115	-7 ^d
Ethylamine	CH ₃ CH ₂ NH ₂	45	8.9 ^b	115/160	
Dimethylamine	CH ₃ NHCH ₃	45	8.2 ^b		
Propylamine	CH ₃ CH ₂ CH ₂ NH ₂	59	8.78 ^b		
2-Propananimine	(CH ₃) ₂ CHNH ₂	59	8.72 ^b		
N-Methyl ethanamine	CH ₃ CH ₂ NHCH ₃	59	8.15 ^f		
N-Methyl-methanediamine	NH ₂ CH ₂ NHCH ₃	60	8.23 ^a		
Ethylenediamine	NH ₂ CH ₂ CH ₂ NH ₂	60	8.27 ^a	160	
1,2-Dimethylhydrazine	CH ₃ NHCH ₂ CH ₃	60	7.18 ^a		
1,1-Dimethylhydrazine	(CH ₃) ₂ NNH ₂	60	7.24 ^a		
Protonated methylamine	CH ₃ NH ₃ ⁺	32			
Methylamine isotope	¹³ CH ₃ NH ₂	32			
Protonated methylamine isotope	¹³ CH ₃ NH ₃ ⁺	33			
Photofragments for ethylenediamine and methylamine					
Methylamine	CNH ₄ ⁺	30	AE = 10.18 ^e	115	
Ethylenediamine	CNH ₄ ⁺	30	AE = 9.3 ^c	160	
Ethylenediamine	C ₂ NH ₅ ⁺	43	AE = 8.85 ^c	160	
Ethylenediamine	C ₂ NH ₆ ⁺	44	AE = 8.90 ^c	160	
Ethylenediamine	C ₂ N ₂ H ₆ ⁺	58 ^c		160	
Ethylenediamine	C ₂ N ₂ H ₇ ⁺	59	AE = 9.06 ^c	160	

^a This work. ^b Taken from Watanabe *et al.*, 1957.⁷⁸ ^c Taken from Wei *et al.*, 2006.⁸⁰ ^d Taken from Ruscic *et al.*, 2016.⁹⁹ ^e Taken from Lossing *et al.*, 1987.¹⁰⁰ ^f Taken from Aue *et al.*, 1979.¹⁰¹

energy being supplied from the ionizing radiation *via* inelastic energy transfer processes. Previous infrared studies after Photolysis by Bossa *et al.*³⁴ indicated that both formation routes might lead to CH₂NH. These mechanisms were also documented in gas phase studies by Reed *et al.*⁸² utilizing UV photodissociation of CH₃NH₂ at 215 nm in a molecular beam. This work revealed the atomic hydrogen loss pathway and formation of methylamidogen (CH₃NH) (3a), which was enhanced by a factor of three compared to CH₂NH₂ (4a). A summary of the reaction pathways discussed in this study are displayed in Fig. 4.

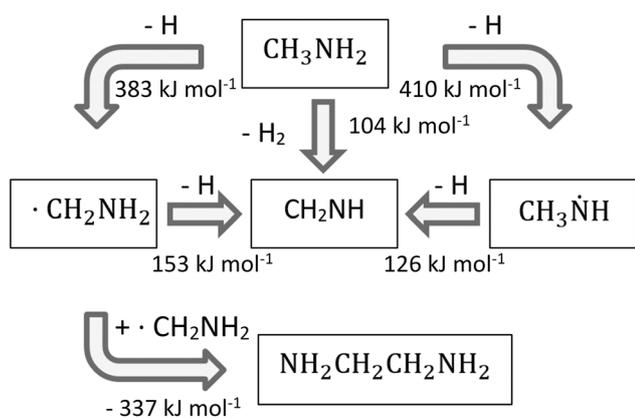


Fig. 4 Summary of reaction pathways toward CH₂NH and NH₂CH₂CH₂NH₂. The Gibbs free energy change of each reaction is also indicated.

A possible formation pathway for the NH₂CH₂CH₂NH₂ molecule could follow a barrier-less radical-radical reaction *via* the exoergic recombination of two CH₂NH₂ radicals (reaction (5)). At low temperatures, these radical-radical recombination pathways are very prominent in interstellar ices, as long as the radicals are next to each other and the recombination geometry is correct. This has been successfully demonstrated for example for the formation of ethane (CH₃CH₃),^{15,83,84} hydrazine (NH₂NH₂),⁸⁵ diphosphine (PH₂PH₂),⁴⁸ hydrogen peroxide (HOOH),^{86,87} ethylene glycol (HOCH₂CH₂OH),⁴¹ and hydroxylamine (NH₂OH),⁵⁰ *via* recombination of methyl (CH₃), amido (NH₂), phosphino (PH₂), hydroxyl (OH), and amido (NH₂) with hydroxyl (OH).

Ogura *et al.*⁸⁸ suggested a pathway in analogy to reaction (5). In their gas phase study, they photolyzed gas mixtures of CH₄, NH₃ and H₂O with broadband UV with the highest fluxes at 185 and 254 nm. They detected NH₂CH₂CH₂NH₂ among others. It should be stressed that the overall reactions to form CH₂NH and NH₂CH₂CH₂NH₂ from CH₃NH₂ are endoergic by 536 kJ mol⁻¹ (reaction (6)) and 463 kJ mol⁻¹ (reaction (7)), respectively. Therefore, these reactions cannot happen thermally at 10 K or during the warm up phase, but have to involve non-equilibrium chemistry.⁴⁰ Even a hypothetical molecular hydrogen elimination (reaction (8)) is still endoergic by 104 kJ mol⁻¹. Therefore, the reaction pathways involved demand for the presence of an external energy source provided by, *e.g.*, secondary electrons generated by GCRs in order to successfully form the product in the ISM.

Table 5 Comparison of CH₃NH₂ and CH₃OH reaction pathways. Data were taken from the NIST Chemistry WebBook and active thermochemical tables

Reaction number	Reaction	$\Delta_R G$
CH ₃ NH ₂		
3a	CH ₃ NH ₂ → CH ₃ NH + H	410 kJ mol ⁻¹ ; 4.25 eV
3b	CH ₃ NH → CH ₂ NH + H	126 kJ mol ⁻¹ ; 1.31 eV
4a	CH ₃ NH ₂ → CH ₂ NH ₂ + H	383 kJ mol ⁻¹ ; 3.97 eV
4b	CH ₂ NH ₂ → CH ₂ NH + H	153 kJ mol ⁻¹ ; 1.59 eV
5	CH ₂ NH ₂ + CH ₂ NH ₂ → NH ₂ CH ₂ CH ₂ NH ₂	-337 kJ mol ⁻¹ ; -3.50 eV
6	CH ₃ NH ₂ → CH ₂ NH + 2H	536 kJ mol ⁻¹ ; 5.56 eV
7	CH ₃ NH ₂ + CH ₃ NH ₂ → NH ₂ CH ₂ CH ₂ NH ₂ + 2H	463 kJ mol ⁻¹ ; 4.80 eV
8	CH ₃ NH ₂ → CH ₂ NH + H ₂	104 kJ mol ⁻¹ ; 1.08 eV
CH ₃ OH		
9a	CH ₃ OH → CH ₃ O + H	435 kJ mol ⁻¹ ; 4.51 eV
9b	CH ₃ O → CH ₂ O + H	82 kJ mol ⁻¹ ; 0.85 eV
10a	CH ₃ OH → CH ₂ OH + H	396 kJ mol ⁻¹ ; 4.10 eV
10b	CH ₂ OH → CH ₂ O + H	121 kJ mol ⁻¹ ; 1.25 eV
11	CH ₂ OH + CH ₂ OH → HOCH ₂ CH ₂ OH	-350 kJ mol ⁻¹ ; -3.63 eV
12	CH ₃ OH → CH ₂ O + 2H	517 kJ mol ⁻¹ ; 5.36 eV
13	CH ₃ OH + CH ₃ OH → HOCH ₂ CH ₂ OH + 2H	449 kJ mol ⁻¹ ; 4.65 eV
14	CH ₃ OH → CH ₂ O + H ₂	85 kJ mol ⁻¹ ; 0.88 eV

6. Conclusions

The present study explored the formation of CH₂NH *via* the unimolecular decomposition of CH₃NH₂ in interstellar ices. Methylamine is present in several interstellar clouds such as Sagittarius B2²³ and Orion-KL^{1,24} and expected to be incorporated in interstellar ices as well. Even though it is expected that CH₃NH₂ is present only as a minor component in the ice, the present experiments reveal that each CH₂NH product can result from fragmentation of one single CH₃NH₂ molecule. These results of our mechanistical studies are transferable to interstellar ice mixtures containing CH₃NH₂, which are exposed to ionizing radiation. The irradiation dose utilized in this study is equivalent to 10⁵ years of the life time of a cold molecular cloud.⁸⁹ Here, CH₂NH represents an important intermediate to the abiotic formation of extraterrestrial amino acids in interstellar ices,¹⁴ which are essential building blocks for life on Earth.⁹⁰ The data collected by the elegant PI-ReTOF-MS approach highlight the feasibility of next generation laboratory studies to detect radiation and reaction products of non-equilibrium chemistries in interstellar analog ices isomers selectively. In addition to the formation of CH₂NH, the experiments confirmed the formation of NH₂CH₂CH₂NH₂ in the irradiated CH₃NH₂ ice. This has not been observed in the interstellar medium up to date. A possible explanation might be found by the lower fractional abundance of the parent molecule CH₃NH₂ in the interstellar ices compared to our analog ice samples. This delimits CH₃NH₂ molecules present in direct neighborhood to each other in the ice and thus the recombination of two subsequently formed CH₂NH₂ radicals *via* radical-radical reaction is delimited as well. As presented earlier, methylamine and CH₂NH are present in several molecular clouds and star forming regions as well as the dust samples collected by the Stardust mission⁴ and the comet 67P/Churyumov-Gerasimenko.²⁹ Ethylenediamine has not been observed in the interstellar medium or in the solar system yet, but our results suggest that it should be detectable in those

environments where CH₃NH₂ and CH₂NH have been discovered. The upper limits of CH₂NH and of NH₂CH₂CH₂NH₂ in this study are below 0.09 ± 0.01 and 0.045 ± 0.005 molecules eV⁻¹, respectively, and these formation efficiency can be utilized in future astrochemical models of non-equilibrium ice chemistry to predict its interstellar abundance.⁹¹

It is interesting to note that CH₃NH₂ is isoelectronic to CH₃OH. Methanol can reach fractions of up to 27% with respect to H₂O in interstellar ices. Maity *et al.*^{41,45-47} irradiated CH₃OH bearing ices at 5–10 K by energetic electrons. Here, methanol, which consists of a methyl (CH₃) group and a hydroxyl (OH) group (Table 5), has been found to undergo similar reaction routes compared to CH₃NH₂, which in turn carries a methyl (CH₃) group and an amine (NH₂) group. In the CH₃OH ice, the product H₂CO – isoelectronic to CH₂NH – was formed *via* either a methoxy radical (CH₃O) or a hydroxymethyl radical intermediate (CH₂OH) *via* unimolecular decomposition pathways (reactions (9) and (10), Table 5). These reactions are in analogy to pathways (3) and (4) in the present study and are endoergic as well. Each pair of analogous reactions has similar Gibbs free energy changes, which may induce the observed same trends of column densities of N_{CH₃NH₂} > N_{CH₂NH} and N_{CH₃OH} > N_{H₂CO} in Sagittarius B2.⁹²⁻⁹⁴ Likewise, ethyleneglycol (HOCH₂CH₂OH) – isoelectronic to NH₂CH₂CH₂NH₂ – has been found to be synthesized from the hydroxymethyl (CH₂OH) *via* radical-radical reaction in CH₃OH ices and has been detected in the interstellar medium toward Sagittarius B2(N-LMH).⁹⁵ This radical-radical formation is similar to the formation of NH₂CH₂CH₂NH₂ (5) *via* the CH₂NH₂ in CH₃NH₂ ice in this study. Therefore, ethylenediamine is expected to be present in these interstellar environments, where CH₃NH₂ and CH₂NH has been observed as well such as towards Sagittarius B2.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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