



Methane Formation in Cold Regions from Carbon Atoms and Molecular Hydrogen

Thanja Lamberts^{1,2} , Gleb Fedoseev^{2,3} , Marc C. van Hemert¹, Danna Qasim^{2,4} , Ko-Ju Chuang² , Julia C. Santos² , and Harold Linnartz²

¹ Leiden Institute of Chemistry, Gorlaeus Laboratories, Leiden University, P.O. Box 9502, 2300 RA Leiden, The Netherlands; a.l.m.lamberts@lic.leidenuniv.nl

² Laboratory for Astrophysics, Leiden Observatory, Leiden University, P.O. Box 9513, 2300 RA Leiden, The Netherlands

³ Research Laboratory for Astrochemistry, Ural Federal University, Kuibysheva St. 48, 620026 Yekaterinburg, Russia

Received 2021 September 16; revised 2022 January 18; accepted 2022 February 2; published 2022 March 25

Abstract

Methane is typically thought to be formed in the solid state on top of cold interstellar icy grain mantles via the successive atomic hydrogenation of a carbon atom. In the current work we investigate the role of molecular hydrogen in the CH₄ reaction network. We make use of an ultrahigh vacuum cryogenic setup combining an atomic carbon atom beam with atomic and/or molecular beams of hydrogen and deuterium on a water ice. These experiments lead to the formation of methane isotopologues detected in situ through reflection absorption infrared spectroscopy. Most notably, CH₄ is experimentally formed by combining C atoms with only H₂ on amorphous solid water, albeit more slowly than in experiments where H atoms are also present. Furthermore, CH₂D₂ is detected in an experiment involving C atoms with H₂ and D₂ on H₂O ice. CD₄, however, is only formed when D atoms are present in the experiment. These findings have been rationalized by means of computational and theoretical chemical insights. This leads to the following conclusions: (a) the reaction C + H₂ → CH₂ takes place, although it is not barrierless for all binding sites on water, (b) the reaction CH + H₂ → CH₃ is barrierless, but has not yet been included in astrochemical models, (c) the reactions CH₂ + H₂ → CH₃ + H and CH₃ + H₂ → CH₄ + H can take place only via a tunneling mechanism, and (d) molecular hydrogen possibly plays a more important role in the solid-state formation of methane than assumed so far.

Unified Astronomy Thesaurus concepts: Astrochemistry (75); Interstellar molecules (849); Molecular spectroscopy (2095); Molecular clouds (1072); Laboratory astrophysics (2004); Computational methods (1965)

1. Introduction

Methane, the smallest hydrocarbon, is one of the few molecules that have been detected in the solid phase in various regions in the interstellar medium (Boogert et al. 2015). In fact, the first detection was a simultaneous gas-phase and tentative solid-phase identification, based on the ν_4 feature at 7.6 μm (Lacy et al. 1991); meanwhile several in-depth observational studies have been reported (Boogert et al. 1996; Öberg et al. 2008). Early reports based on comparison to laboratory data indicated that methane likely resides in ice comprising polar component(s) (Boogert et al. 1996), and it was later postulated that H₂O is the primary candidate for this based on correlations between CH₄ and H₂O column densities (Öberg et al. 2008). This points to the fact that solid CH₄ is formed during the translucent phase of the evolutionary track of molecular clouds.

The solid-state formation of methane has typically been assumed to follow four sequential atomic hydrogenation steps of the carbon atom in the ³P ground state ever since this was postulated in the late 1940s (van de Hulst 1946, 1949; D'Hendecourt et al. 1985; Brown et al. 1988; Brown & Charnley 1991). Recently, this route has been confirmed experimentally (Qasim et al. 2020a) through the simultaneous use of well-characterized C- and H-atom beams, following up

on early work by Hiraoka et al. (1998):



The hydrogen atom number density in molecular clouds is estimated to be around a few atoms cm⁻³, which is two to four orders of magnitude lower than the molecular hydrogen abundance depending on whether a translucent or dense cloud is concerned (van Dishoeck & Black 1988; Goldsmith & Li 2005). Therefore, solid-state reactions with molecular hydrogen can also be of great importance even when the corresponding rate constants are lower, as pointed out already by Hasegawa & Herbst (1993). For instance, for the sequential hydrogenation of the O atom to eventually form water (Hiraoka et al. 1998; Ioppolo et al. 2008; Miyauchi et al. 2008) it has been shown that the reaction H₂ + OH → H₂O + H can become more relevant than H + OH → H₂O even though a considerable barrier is invoked (Cuppen & Herbst 2007; Furuya et al. 2015). Another example is the hydrogenation of carbon monoxide by UV irradiation of mixed CO:H₂ ices (Chuang et al. 2018). Despite such active involvement of H₂ in solid-state reactions, the molecular hydrogen abundances on ice surfaces in astrochemical microscopic models are often artificially reduced to save (a lot of) computational cost, e.g., by decreasing the sticking coefficient (Garrod 2013; Lamberts et al. 2013; Vasyunin & Herbst 2013). We also want to point out that, despite the fact that neutral carbon atoms are often thought to only be (abundantly) present in translucent regions (van Dishoeck & Black 1988; Snow & McCall 2006), there

⁴ Current address: Astrochemistry Laboratory, NASA Goddard Space Flight Center, Greenbelt, MD 20771, USA.

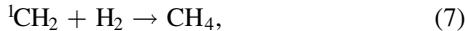
Original content from this work may be used under the terms of the [Creative Commons Attribution 4.0 licence](#). Any further distribution of this work must maintain attribution to the author(s) and the title of the work, journal citation and DOI.

exists substantial literature that indicates that C [I] is more extended, possibly even into denser regions (Langer 1976; Keene et al. 1985; Papadopoulos et al. 2004; Burton et al. 2015; Bisbas et al. 2019).

For these reasons, we consider a number of reactions with molecular hydrogen in the context of methane formation. First, the direct addition or insertion reactions:

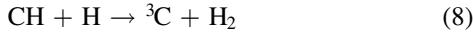


Reaction (5), the first step in the reaction network, has been covered in a series of papers indicating that the reaction may readily take place in helium droplets (Krasnokutski et al. 2016; Henning & Krasnokutski 2019), and including this reaction in astrochemical models without a barrier was suggested recently (Simončič et al. 2020). The reaction



intuitively the most likely step to form methane, can only take place if methylene is in the excited singlet state, ${}^1\text{CH}_2$ (Murrell et al. 1973; Bauschlicher et al. 1977). In the system currently under study experimentally, only ground-state ${}^3\text{CH}_2$ is expected to be present, and therefore reaction (7) will not be further considered. Excited-state singlet carbon atoms can, however, play a role in other interstellar environments, for instance Titan’s atmosphere (Hickson et al. 2016), but this is beyond the scope of the current paper.

In terms of the chemical reaction network, there is furthermore the possibility of hydrogen abstraction, either from H_2 or from a CH_n fragment:



Reactions (8)–(11) are listed in the exothermic direction, and the reverse endothermic reactions are not expected to be important given the low temperatures involved (~ 10 –20 K). We assume that the exothermicities of the reactions are not significantly altered by the solid-state surroundings of the reaction site. This essentially translates into assuming that the binding energies of the reactants and products are similar. We also assume that the H atoms formed in reactions (10) and (11) immediately desorb, based on the argument of conservation of energy and momentum (Koning et al. 2013).

In the current paper, we revisit reaction (5) in water ices and extend the discussion on H_2 reactivity, investigating the influence of reactions (6)–(11). We show that reactions (5), (10), and (11) together can lead to the formation of methane, without the involvement of H atoms. To achieve this, we make use of in situ infrared spectroscopy to probe which methane isotopologues are formed via reaction of ${}^3\text{C}$ with selected combinations of atomic and molecular hydrogen (H , H_2) and deuterium (D , D_2) on representative interstellar water-rich ice analogs.

This paper is organized in the following way. The experimental details and results are discussed in Sections 2.1 and 3. Furthermore, we position the chemical network listed above within the context of the extensive theoretical chemical

literature, complemented by additional computations presented that explicitly take into account the role of the water ice surface. This helps to disentangle which reactions are likely to take place throughout the various experiments, as outlined in Section 2.2 and discussed in Section 4. Solid methane is hard to observe from ground-based observatories because of telluric pollution. Observations from space offer an alternative. Solid methane has been observed already with the Spitzer space telescope (Öberg et al. 2008). Because of its higher sensitivity and spatial resolution, the James Webb Space Telescope (JWST) is expected to substantially extend these observations. The present experimental work and theoretical approach fits worldwide efforts to prepare for upcoming JWST observations. In Section 5 the astrochemical implications and conclusions of this work are presented.

2. Methodology

2.1. Experimental Methodology

The experimental setup used is SURFRESIDE³, an ultrahigh vacuum (UHV) system with three atomic beam lines (Ioppolo et al. 2013; Qasim et al. 2020b). For the purpose of our study, only H/H_2 , D/D_2 , and C atom beam lines are used. Ices are grown on a gold-coated copper substrate that is attached to the cold finger of a closed-cycle He cryostat in the center of the main UHV chamber with a base pressure of the order of 10^{-10} mbar. Co-deposition experiments of $\text{H}_2\text{O} + ({}^3\text{P})\text{C}$ and combinations of H, D, H_2 , and/or D_2 are performed that lead to the growth of a mixed ice at 10 K. Mixed H/H_2 or D/D_2 beams are obtained by (partial) dissociation of molecular H_2 (Linde 5.0) or D_2 (Linde 2.8) in a microwave discharge atom source (MWAS, Oxford Scientific, Schmidt et al. 1996; Anton et al. 2000) in a separate vacuum chamber with a base pressure of $\sim 10^{-9}$ mbar. Note that charged particles are removed by applying an electric field that deflects these species. Excited-state species are de-excited through collisions with the walls of a U-shaped quartz pipe at room temperature placed along the beam path prior to the molecules entering the main chamber. A customized SUKO-A 40 C-atom source (Qasim et al. 2020b) based on a commercial design (Dr. Eberl, MBE, Krasnokutski & Huisken 2014; Albar et al. 2017) produces a beam of carbon atoms in the ${}^3\text{P}$ ground state, albeit with a mean translational kinetic energy of the order of the source temperature (~ 2000 K), with a C_n/C ($n > 1$) ratio of less than 0.01. Based on recent calculations for the energy dissipation timescale of less than a few picoseconds for translationally hot nitrogen atoms on amorphous solid water, we expect no impact of the high carbon source temperature on the possible chemistry that takes place (Molpeceres et al. 2020). This source is located in another separate vacuum chamber with a base pressure $(3\text{--}5) \times 10^{-9}$ mbar. A series of apertures is used to collimate the C-atom beam on the substrate, avoiding deposition of carbon on the walls of the main UHV chamber. Note that CO and CO_2 are known contaminants in the C-atom beam. A third vacuum chamber with a base pressure of $\sim 10^{-9}$ mbar is used to generate a molecular H_2O (Milli-Q) or D_2O (Sigma-Aldrich, 99.9 at.% D) beam for simultaneous deposition (molecular deposition line, MDL). We use an overabundance of H_2O to mimic a polar ice environment, e.g., in agreement with the previously mentioned correlation between observed CH_4 and H_2O column densities. Water reaches the surface with a translational kinetic energy equal to or less than room

Table 1
List of Experimental Conditions (Temperature, Flux, Mixing Ratio and Time), Organized into Four Selected Sets

| # | T (K) | H ₂ O (cm ⁻² s ⁻¹) | C (cm ⁻² s ⁻¹) | H (cm ⁻² s ⁻¹) | H ₂ (cm ⁻² s ⁻¹) | D (cm ⁻² s ⁻¹) | D ₂ (cm ⁻² s ⁻¹) | C:H:H ₂ :D:D ₂ | Time (min) |
|-----|-----------|---|--|---|---|--|---|--------------------------------------|---------------|
| | | MDL | SUKO | MWAS | MWAS | | MDL | | |
| 1 A | 10 | 8×10^{12} | 5×10^{11} | 2×10^{12} | 1×10^{14} | | | 1:4:200:-:- | 30 |
| 1 B | 10 | 8×10^{12} | 5×10^{11} | 2×10^{12} | 1×10^{14} | | 1×10^{13} | 1:4:200:-:20 | 30 |
| 1 C | 10 | 8×10^{12} | 5×10^{11} | 2×10^{12} | 1×10^{14} | | 4×10^{13} | 1:4:200:-:80 | 30 |
| | | MDL | SUKO | | MDL | MWAS | MWAS | | |
| 2 A | 10 | 8×10^{12} | 5×10^{11} | | | 1.5×10^{12} | 6×10^{13} | 1:-:-:3:120 | 30 |
| 2 B | 10 | 8×10^{12} | 5×10^{11} | | 5×10^{13} | 1.5×10^{12} | 6×10^{13} | 1:-:100:3:120 | 30 |
| 2 C | 10 | 8×10^{12} | 5×10^{11} | | 2×10^{14} | 1.5×10^{12} | 6×10^{13} | 1:-:400:3:120 | 60 |
| | | MDL | SUKO | | MDL | | MDL | | |
| 3 A | 10 | 1.2×10^{13} | 5×10^{11} | | | | 1×10^{14} | 1:-:-:-:200 | 60 |
| 3 B | 10 | 1.2×10^{13} | 5×10^{11} | | 1×10^{14} | | 1×10^{14} | 1:-:200:-:200 | 240 |
| # | T (K) | D ₂ O (cm ⁻² s ⁻¹) | C (cm ⁻² s ⁻¹) | H ₂ (cm ⁻² s ⁻¹) | | | | | Time (min) |
| | | MDL | SUKO | | MDL | | | | |
| 4 A | 10 | 1.4×10^{13} | 5×10^{11} | | 2.5×10^{14} | | | 1:-:500:-:- | 60 |
| 4 B | 25 | 1.4×10^{13} | 5×10^{11} | | 2.5×10^{14} | | | 1:-:500:-:- | 60 |

Note. MDL, SUKO, and MWAS refer to the atomic and molecular deposition lines used (see text). Furthermore the substrate temperature, the atomic and molecular fluxes, and the total time of the experiment, from which the fluence can be derived, are listed. All fluxes give the effective value. Note that H (H₂) and D (D₂) have different thermal velocities and sticking coefficients, thus a direct comparison should be made with care. Values in bold font underline the different settings within one series of measurements.

temperature. Based on the lack of dangling modes in the infrared spectra, we conclude that the co-deposition of C, H/H₂, and H₂O does not lead to the deposition of a very porous amorphous solid water ice. The atomic and molecular fluxes are listed in Table 1, in which all experiments are summarized. The uncertainty in the flux determination is roughly a factor of two. The initial reactants and formed products are monitored in situ via reflection absorption infrared spectroscopy (RAIRS, Greenler 1966) in the solid state. Note that the infrared beam is guided toward and from the entrance and exit windows of the chamber via boxes that are purged with dry air to reduce atmospheric absorption. Finally, a baseline correction is performed that includes the subtraction of known components of the purging gas used along the path of the infrared beam, i.e., outside the vacuum chamber, such as water vapor. The experiments are organized into four different series in which relevant parameters are systematically varied—see Table 1.

2.2. Computational Methodology

The primary focus of our own calculations is on the reaction C + H₂ → CH₂, since it is the first and most determining step in the reaction network mentioned above leading to CH₄. Moreover, the carbon atom is known to be highly reactive (Kim et al. 2003). However, with the exception of the work by Simončič et al. (2020), to date it has not been looked at it in detail for surface chemistry purposes. Note that for the other possible steps (reactions (6)–(11)), we draw from previous studies available from the literature.

We take a threefold approach—see also Figure 1—to understand the reactivity of C + H₂ → CH₂:

1. a gas-phase calculation of a highly symmetric orientation of C with respect to H₂ (C_{2v} symmetry)—meant to compare against previous results,
2. a gas-phase calculation of a low-symmetry orientation of C with respect to H₂ (C_s symmetry)—meant to be a first step toward a realistic symmetry-broken orientation on a surface,
3. the calculation of the reaction of H₂ with a C atom bound to a single H₂O molecule and an (H₂O)₃ cluster (no symmetry)—meant as a first step toward understanding the reaction with a bound carbon atom on a water ice.

When a gas-phase H₂ molecule approaches a carbon atom in its ³P ground state the encounter can occur on three, initially degenerate, potential surfaces (PESs). Important to take into account is the alignment of the two singly occupied 2p orbitals of the carbon atom. When the orientation of the H₂ molecule is perpendicular to the direction of approach, there is a C_{2v} symmetry and the three surfaces are ³B₁, ³B₂, and ³A₂. For all three surfaces the energy was calculated as a function of the Jacobi coordinates *R* as the distance between C and the center of mass of H₂, *r* as the H–H distance, and $\theta = 90^\circ$ as the angle between *R* and *r*; see again Figure 1. All calculations were repeated in C_s symmetry, where the angle θ was reduced to 80° , while keeping the CAS/CI parameters as close as possible to the ones used in the C_{2v} case. In C_s symmetry two ³A'' states are relevant. Subsequently, to find possible reaction paths, the location of the crossing (seam) between the ³B₁ and ³A₂ for C_{2v} and the two ³A'' surfaces for C_s was searched for by scanning these surfaces with small steps in the $R \approx r \approx 1 \text{ \AA}$ range. These calculations have been performed using Molpro (Werner et al. 2012, 2018) with the AVQZ basis set (Woon & Dunning 1993; Dunning et al. 2001). The CASSCF calculation had eight

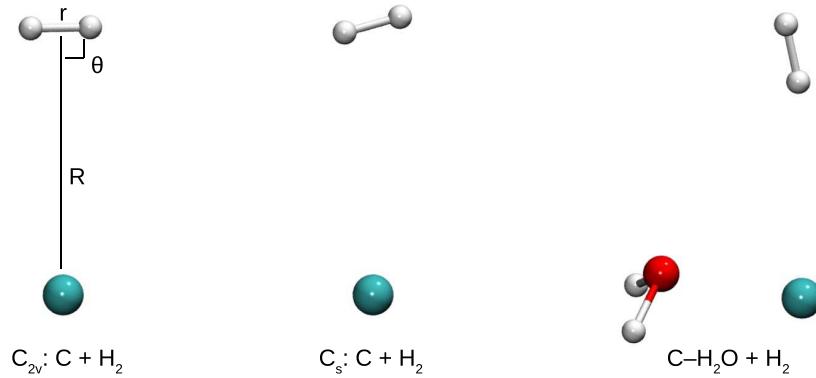


Figure 1. Three cases studied for the reaction $\text{C} + \text{H}_2 \rightarrow \text{CH}_2$; the reaction on the $(\text{H}_2\text{O})_3$ cluster is not depicted here.

electrons in eight orbitals, while state-averaging over singlet and triplet states, each with four lowest roots, for both C_{2v} and C_s symmetries.

In order to test—for the first time—the influence of the strong adsorption of the carbon atom atop water ice (see also Molpeceres et al. 2021) on the reactivity of the carbon atom with H_2 we have performed a nudged elastic band calculation for the reaction of H_2 with a carbon atom bound to a water molecule, i.e., a $\text{C}-\text{H}_2\text{O}$ complex, and to a water trimer $(\text{H}_2\text{O})_3$, i.e., a $\text{C}-(\text{H}_2\text{O})_3$ complex. This was followed by a transition state optimization applying the dimer method with the B3LYP functional (Becke 1988, 1993; Lee et al. 1988) and a def2-TZVP basis set (Weigend & Ahlrichs 2005), using DL-find in Chemshell (Kästner et al. 2009; Metz et al. 2014). For the calculations with the single water molecule, single-point energies were calculated for the reactant, transition, and product states, with both MRCI/AVTZ (Werner & Knowles 1988; Dunning 1989; Kendall et al. 1992) and CCSD(T)-F12a/VTZ-F12 (Adler et al. 2007; Peterson et al. 2008; Knizia et al. 2009) in Molpro version 2020 (Werner et al. 2020). All geometries were reoptimized at the CCSD(T)-F12a/VTZ-F12 level of theory in Molpro. MRCI calculations are, in principle, warranted for systems where multiple reference effects can be expected, such as here where a triplet ground state carbon atom is involved. We do note, however, that the $T1$ and $D1$ values in both CCSD(T)-F12 calculations are (well) below the common threshold values of $T1 < 0.04$ and $D1 < 0.05$ (Janssen & Nielsen 1998; Lambert et al. 2006). Indeed, the MRCI single-point energy calculations indicate that the main contribution of a Slater determinant to the wave function has reference coefficients of about 0.93–0.94 for all $\text{C}-\text{H}_2\text{O}-\text{H}_2$ geometries.

Note that, for the water monomer, all higher level calculations are in good agreement with the energetics predicted by B3LYP/def2-TZVP (see Table 2), and therefore the results for the water trimer are expected to be representative.

3. Experimental Results

Figure 2 shows the RAIR spectra of experimental series 1 and 2, over the wavenumber ranges that include all relevant spectroscopic bands of CH_4 , CD_4 , and CH_2D_2 , indicated by the vertical lines in the figure. Other detected species are listed in Table 4 and discussed in Appendix B.

Table 2
Activation and Reaction Energies in kJ mol^{-1} , E_{act} and E_{react} Respectively, for the Reaction of the $\text{C}-(\text{H}_2\text{O})_n$ Complex with H_2 Leading to the $\text{CH}_2-(\text{H}_2\text{O})_n$ Complex, with Respect to the Pre-reactive Complex

| | MRCI/ B3LYP | CCSD(T)-F12/ B3LYP | CCSD (T)-F12 |
|---|----------------|-----------------------|-----------------|
| $E_{\text{int},\text{C}-\text{H}_2\text{O}}$ | −52.9 | | −36.1 |
| $E_{\text{act},\text{H}_2 \dots \text{C}-\text{H}_2\text{O}}$ | 30.4 | 30.0 | 30.4 |
| $E_{\text{react},\text{CH}_2-\text{H}_2\text{O}}$ | −294.3 | −281.5 | −300.5 |
| $E_{\text{int},\text{CH}_2-\text{H}_2\text{O}}$ | −8.0 | | −7.1 |
| $E_{\text{int},\text{C}-(\text{H}_2\text{O})_3}$ | −95.2 | | |
| $E_{\text{act},\text{H}_2 \dots \text{C}-(\text{H}_2\text{O})_3}$ | 77.9 | | |
| $E_{\text{react},\text{CH}_2-(\text{H}_2\text{O})_3}$ | −250.9 | | |
| $E_{\text{int},\text{CH}_2-(\text{H}_2\text{O})_3}$ | −9.2 | | |

Note. The interaction energy, E_{int} of the $\text{CH}_2-(\text{H}_2\text{O})_n$ complex is also given, with respect to the separated radical and water cluster. Note that zero-point energies are not included.

Experiments 1A and 2A (top and bottom lines in Figure 2) serve as a control experiment and can be directly compared to previous work (Qasim et al. 2020a). Indeed the formation of CH_4 and CD_4 is clearly confirmed by the presence of both the ν_3 and ν_4 vibrational modes. Molecular deuterium (hydrogen) is introduced in experiment 1B (2B) and increased in experiment 1C (2C), and a concomitant decrease of CH_4 (CD_4) can be easily observed. At the same time, several CH_2D_2 absorption features appear in these four experiments, with the most intense peak at 1028 cm^{-1} . Experiment 1C shows the most apparent, multiline detection of doubly deuterated methane as a result of the advantageous ratio between all reactants: $\text{C:H:H}_2:\text{D}_2 = 1:4:200:80$, see Table 1.

Because doubly deuterated methane is observed in experiments for which either hydrogen or deuterium is present only in the molecular form, at least one reaction with a molecular species must take place throughout the course of methane formation. This is further supported by a tentative CH_4 detection at 1303 cm^{-1} in experiment 2C, i.e., in an experiment with hydrogen present only in the molecular form.

In fact, this tentative detection was the reason to perform experimental series 3 and 4 to further investigate whether methane indeed can be formed if only molecular forms of

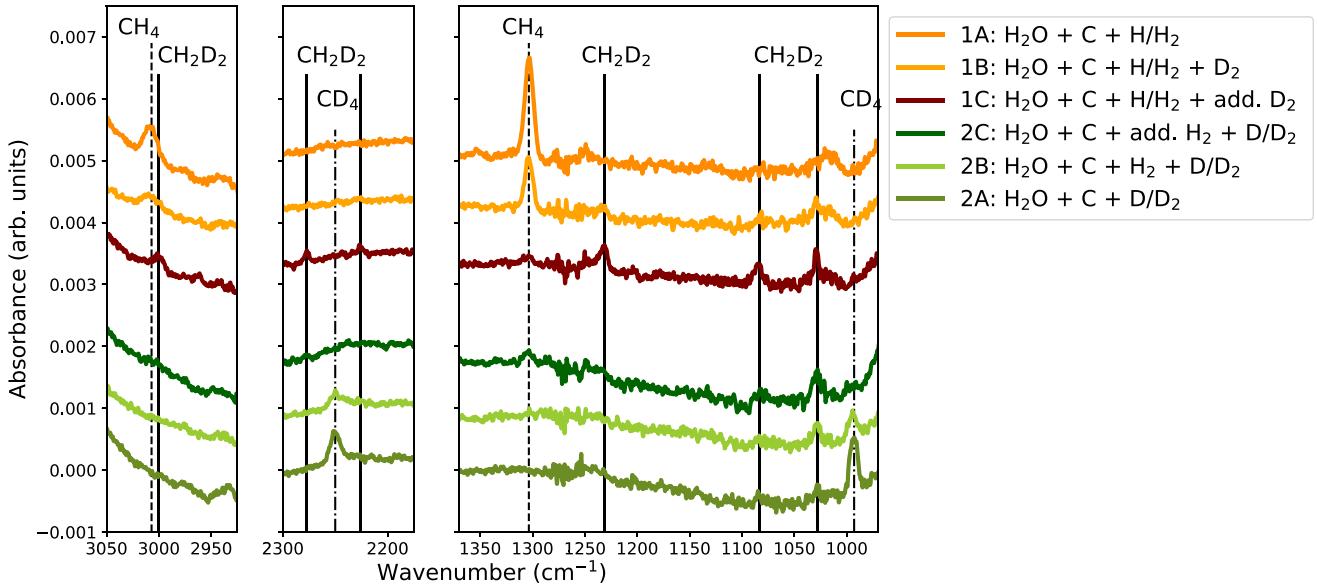


Figure 2. RAIR spectra of experimental series 1 and 2; the exact experimental conditions can be found in Table 1. The dashed vertical lines indicate the peak positions of CH_4 , the solid vertical lines indicate CH_2D_2 , and the dashed-dotted lines indicate CD_4 ; see Table 4 in Appendix B for specific peak positions.

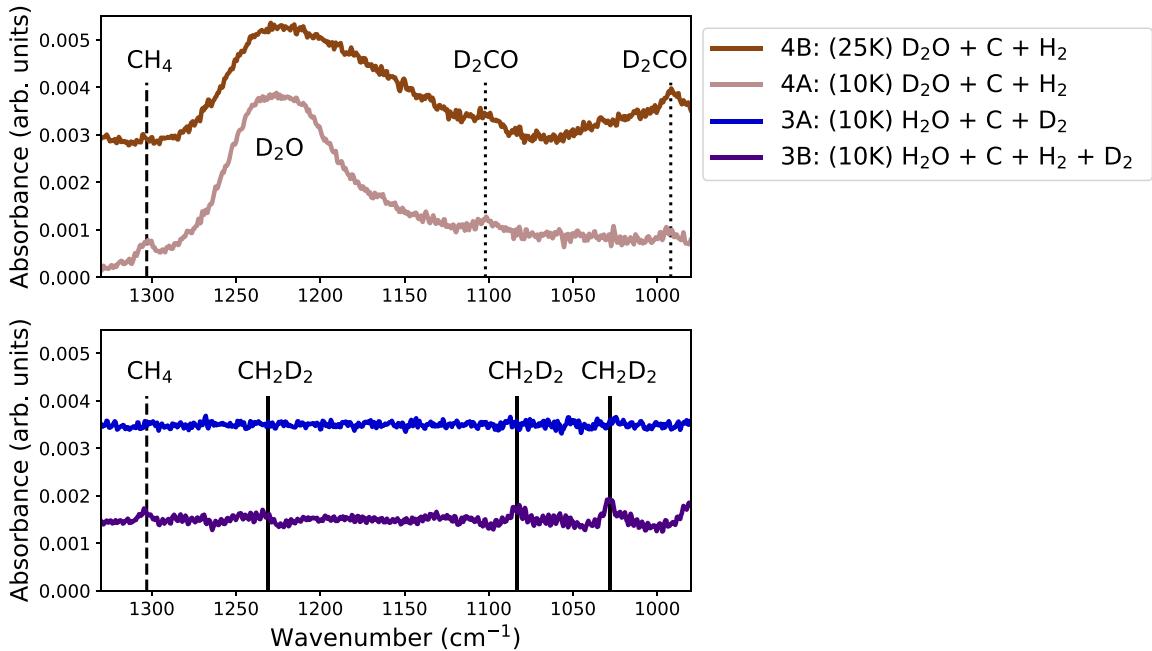


Figure 3. RAIR spectra of experiments 3A, 3B, 4A, and 4B; the exact experimental conditions can be found in Table 1. The dashed vertical lines indicate the peak position of CH_4 , the solid vertical lines indicate CH_2D_2 , the dotted lines indicate D_2CO , and the broad peak is caused by D_2O indicated directly in the plot; see Table 4 in Appendix B for specific peak positions.

hydrogen or deuterium are present. In order to increase the signal-to-noise ratio, these experiments have been run over longer times, as indicated in Table 1. The resulting thick ices (~ 200 ML) and interference patterns over the full range of the IR spectra have led to the presentation of baseline-corrected spectra of a small region of interest only ($1330\text{--}980\text{ cm}^{-1}$), depicted in Figure 3.

Experiments 3B, 4A, and 4B clearly result in a lower overall formation rate of methane in comparison to interactions with atomic species. Experiments 3A and 4A represent a

co-deposition of $\text{C} + \text{D}_2 + \text{H}_2\text{O}$, and $\text{C} + \text{H}_2 + \text{D}_2\text{O}$, respectively. In experiment 3A no CD_4 is detected, while experiment 4A shows a clear solid-state CH_4 detection. In other words, in the reaction network for the formation of methane from carbon atoms and molecular hydrogen there is an isotope effect present, which hints at the importance of tunneling in one or more of the involved reactions. Furthermore, experiment 3B, consisting of $\text{C} + \text{H}_2 + \text{D}_2$ on H_2O ice, shows a clear detection of the two main CH_2D_2 peaks at 1083 and 1028 cm^{-1} . In other words, CH_2D_2 has been formed from carbon atoms and

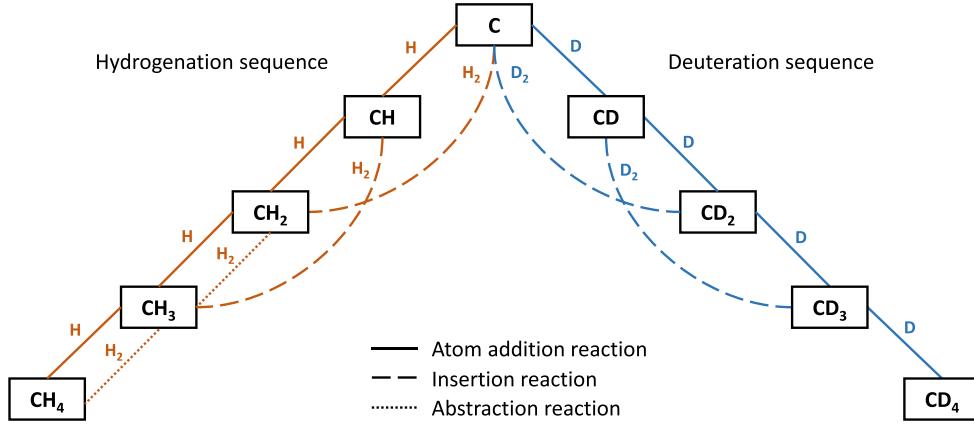


Figure 4. Three types of reactions are considered to lead to the formation of CH₄ or CD₄: H or D addition (Qasim et al. 2020a), H₂ or D₂ insertion (Krasnokutski et al. 2016; Simončič et al. 2020, and this work), and H₂ abstraction (this work). We assume that the latter is not efficient with D₂, based on the involved high barrier and lack of tunneling efficiency.

molecular species only. However, the yield of CH₂D₂ is significantly less than in experiments 1C and 2C.

Experiment 4B, a control experiment, performed at 25 K, i.e., above the desorption temperature of atomic or molecular hydrogen, shows no CH₄ formation in the solid state, confirming that the processes we are studying take place at the surface at low temperatures.

Finally, no CD₃H or CH₃D was detected in any of the experiments, which can be attributed to a combination of isotope effects, low band strengths, and competition between tunneling and barrierless reactions in the network. A full rationale for the lack of these signals is given in Appendix A.

We can summarize the experimental findings as follows.

1. CH₄ is detected in experiments on a water ice using both H atoms and H₂ molecules as well as in experiments using only H₂, albeit with a lower efficiency,
2. CD₄ is detected only in experiments where D atoms are present,
3. CH₂D₂ can be detected in experiments that use simultaneously both atomic and molecular hydrogen or deuterium sources,
4. CH₂D₂ can also be detected in experiments that use exclusively molecular H₂ and D₂,
5. Neither CH₃D nor CHD₃ is detected,
6. All studied reactions take place on the cold ice surface.

4. Theoretical Results

This section focuses on explaining the experimental results by looking in detail at theoretical chemical studies of reactions (5)–(11). Reactions (1)–(4) have been extensively discussed in Qasim et al. (2020a) and will therefore not be further dealt with here. A summary of all relevant reaction steps is depicted in Figure 4.

Section 4.1 contains predominantly results from our own calculations, while Sections 4.2 and 4.3 are based on results available from the literature.

4.1. Reactions on the CH₂ PES

Recently, the reaction C + H₂ → CH₂ was proposed to take place readily on interstellar cold surfaces as well (Simončič et al. 2020). We will first review the reaction in the gas phase and then discuss the impact of having a carbon atom bound to a water ice.

All three calculated gas-phase C_{2v} surfaces for the C + H₂ reaction—³B₁, ³B₂, and ³A₂—are degenerate for large R (C–H₂ distance) with an energy of ∼330 kJ mol⁻¹ above the ³B₁ CH₂ ground state; see Figure 6 in Appendix C. We find a crossing between the ³A₂ and ³B₁ surfaces at R = 1.05 Å and r = 1.17 Å at an energy equal to the asymptotic value of C + H₂. From an energetics point of view and under the assumption that a transition from the ³A₂ to the ³B₁ occurs via a conical intersection, there is no barrier, in agreement with Gamallo et al. (2012). However, at the (avoided) crossing of the C_{2v} potential energy surfaces, there is a change in electron configuration. The dominant contribution to the wave function on the ³A₂ surface consists of an occupation for the six valence electrons of (2a₁)² (3a₁)² (1b₁)¹ (1b₂)¹, and this needs to change to (2a₁)² (3a₁)¹ (1b₁)¹ (1b₂)² before reaching the ³B₁ CH₂ ground state. This means an electron has to move from a p_z orbital on the carbon atom into a p_y orbital, which can impede the reaction from taking place, even if energetically there is no barrier.

When the symmetry is lowered to C_s, which should more closely resemble the situation on an ice surface, by giving up the perpendicular orientation and instead choosing θ = 80°, both the ³B₁ and ³A₂ surfaces become of the same ³A'' symmetry and a reaction can take place without a serious energetic barrier. A small hump can be found along the path at R = 1.08 Å and r = 1 Å, but this remains below the C + H₂ asymptotic value; see Figure 6. Yet the electron configuration still changes upon decreasing R and increasing r, i.e., reacting toward CH₂, and it is unclear what the effect on the reaction efficiency is.

More importantly, however, the known strong interaction of a ³P carbon atom with a water molecule or cluster (Wakelam et al. 2017; Shimonishi et al. 2018; Duflot et al. 2021;

Molpeceres et al. 2021) might drastically change the energetics of the reaction. Table 2 gives an overview of the interaction energy of the C–H₂O complex and the activation and reaction energy of the reaction H₂ + C–H₂O → CH₂–H₂O for different levels of theory. A sizable barrier of 27–30 kJ mol⁻¹ is found, which certainly cannot be trivially overcome at 10 K. For the reaction in the vicinity of the water trimer, the carbon atom is more strongly bound, by ∼42 kJ mol⁻¹, which is directly reflected in a higher barrier of 77 kJ mol⁻¹, i.e., ∼48 kJ mol⁻¹ higher than for the single water molecule. This demonstrates that while the C + H₂ reaction proceeds barrierlessly in helium droplets (Krasnokutski et al. 2016; Henning & Krasnokutski 2019), on water surfaces it will be strongly dependent on the carbon atom binding site. Estimates of the interaction or binding energy depend on the number of water molecules considered and the method used, but generally vary between ∼35 and ∼130 kJ mol⁻¹, much larger than a typical water hydrogen bond of ∼22 kJ mol⁻¹. Our proof-of-principle calculation demonstrates that more extensive work should be done on C–(H₂O)_n clusters with n > 3. At present, therefore, we cannot elaborate on the expected isotope effect of this reaction, but we expect that this reaction on a water ice proceeds with an (effective) barrier and might thus be slower with D₂ than with H₂.

Note that reaction (8), CH + H → ³C + H₂, has to proceed through a CH₂ intermediate, as can be seen in Figure 2 of Gamallo et al. (2012). While this is a realistic scenario in gas-phase experiments in the low-pressure regime where intermediates can convert the reaction energy into internal energy to overcome subsequent barriers, such intermediates are expected to be quenched on a surface by rapid (<1 ps) energy dissipation to a water ice (Fredon et al. 2021). Therefore, reaction (8) turns effectively into reaction (5) on a surface.

4.2. Reactions on the CH₃ PES

The exchange reaction CH + H₂ ⇌ CH₂ + H has been previously studied in the gas phase (McIlroy & Tully 1993; Medvedev et al. 2006; González et al. 2011). In particular, Medvedev et al. (2006) showed that the reaction proceeds through an activated CH₃^{*} intermediate, which is formed via a barrierless pathway in both the forward and backward directions of the reaction, i.e., CH + H₂ ⇌ CH₃^{*} ⇌ CH₂ + H. In the low-pressure regime, González et al. (2011) showed that the three hydrogen atoms become equivalent, as a result of the long-lived CH₃^{*} complex. In the high-pressure regime, on the other hand, McIlroy & Tully (1993) indicated that collisional stabilization to form ground-state CH₃ dominates. A reaction taking place on an ice surface can be seen as an extreme case of the high-pressure limit, in which the ice acts as a third body to take up the excess energy of the reaction (see Section 4.1 above). Given the barrierless nature of the PES, we expect no isotope effect here. In other words, it is highly likely that reaction (6) will lead barrierlessly to the formation of CH₃ on the ice, while reaction (9) is unlikely to take place at all and will revert simply to reaction (3), also forming CH₃. This serves as an explanation why CH₂D₂ is detected in experimental sets 1 and 2, since the barrierless nature leads to a lack of isotope effect.

4.3. Hydrogen Abstraction Reactions

The two remaining reactions with molecular hydrogen, reactions (10) and (11), have been previously studied; the

Table 3
Activation Energies from Predominantly Theoretical Chemical Literature for the Reactions (1)–(6) and (8)–(11) in kJ mol⁻¹

| Reaction | ΔE _{act} | Ref. |
|----------|---|----------------------|
| (1) | 3C + H → CH | 0 [1] |
| (2) | CH + H → CH ₂ | 0 [2] |
| (3) | ³ CH ₂ + H → CH ₃ | 0 [3] |
| (4) | CH ₃ + H → CH ₄ | 0 [4] |
| (5) | ³ C + H ₂ → ³ CH ₂ | TBD ^a [2] |
| (6) | CH + H ₂ → CH ₃ | 0 [3] |
| (8) | CH + H → ³ C + H ₂ | – ^b [2] |
| (9) | CH ₂ + H → CH + H ₂ | – ^c [3] |
| (10) | ³ CH ₂ + H ₂ → CH ₃ + H | 49 [5] |
| (11) | CH ₃ + H ₂ → CH ₄ + H | 44 [6] |

Notes. Note that reaction (7) can only take place in the singlet, excited, state, and is therefore excluded here.

^a Cannot be trivially determined due to conical intersection and strong C–H₂O interaction (see Section 4.1).

^b The reaction is likely quenched in the CH₂ ground state (see Section 4.1 and Gamallo et al. 2012), effectively changing it to reaction (2).

^c Reaction is determined by the relaxation of the CH₃^{*} intermediate (see Section 4.2) and thus leads to reaction (3).

References. [1] Qasim et al. (2020a); [2] Harding et al. (1993); van Harreveld et al. (2002); Gamallo et al. (2012); this work; [3] McIlroy & Tully (1993); Medvedev et al. (2006); González et al. (2011); [4] Duchovic & Hase (1985); [5] Baskin et al. (1974); Bauschlicher (1978); [6] Li et al. (2015); Beyer et al. (2016).

reported barriers are presented in Table 3. Both reactions can only take place when a considerable barrier (>44 kJ mol⁻¹) is overcome, for which tunneling needs to be invoked to reach rate constants that are high enough for the reaction to be able to take place at the low temperatures in dense molecular clouds. The effect of tunneling can be accurately included by means of instanton theory (Miller 1975; Langer 1976; Callan & Coleman 1977; Rommel & Kästner 2011; Richardson 2016). Although Beyer et al. (2016) indeed calculated instanton rate constants for the reaction CH₃ + H₂, they provided only bimolecular rate constants, whereas unimolecular rate constants are those relevant for Langmuir–Hinshelwood-type surface reactions (Lamberts et al. 2016; Meisner et al. 2017). Given the high activation barriers, about twice as large as for the reaction H₂ + OH → H₂O + H (Meisner et al. 2017) and slightly above that of H + H₂O₂ → H₂ + HO₂ (Lamberts et al. 2016), relatively low rate constants are expected. As a consequence of the high barriers, these two reactions are expected to show significant kinetic isotope effects. This, in turn, explains why they are unlikely to take place with molecular deuterium. We interpret the lack of CD₄ detection in our experiments with only D₂ as a deuterium source as experimental confirmation that, indeed, tunneling plays a role in the abstraction reactions from molecular hydrogen and deuterium. A detailed quantitative study will be topic of a future study.

5. Astrochemical Implications and Conclusions

Currently, the main interstellar formation pathway of methane is thought to be through the reactions (1)–(4). Reactions (5), (10), and (11) are included in some astrochemical models, but in most studies the initial guesses by Hasegawa & Herbst are used; see for instance the current surface reactions in the KIDA database (Wakelam et al. 2015). Reaction (6), CH + H₂ → CH₃, is usually not included at all.

For the interpretation of astronomical data, specifically those obtained by the JWST in the near future, it is important for models to take into account that:

1. the reaction $C + H_2 \rightarrow CH_2$ is unlikely to proceed via a fully barrierless mechanism on water ices and an isotope effect is yet to be determined;
2. the reaction $CH + H_2 \rightarrow CH_3$, on the other hand, is expected to take place readily and barrierlessly without an isotope effect;
3. the abstraction reactions $CH_2 + H_2 \rightarrow CH_3 + H$ and $CH_3 + H_2 \rightarrow CH_4 + H$ take place via a tunneling mechanism and a pronounced isotope effect is expected.

The points above are summarized in Table 3 and Figure 4.

The main finding is that H_2 plays a role in the solid-state formation of interstellar methane. CH_4 can be formed without invoking any H atoms in our experiments, which is supported by our own calculations as well as theoretical results found in physical chemical literature. Thus, under physical conditions where H_2 is much more abundant than H atoms and taking into account that H_2 sticks to the surface at higher temperatures than H, methane formation from C atoms and H_2 (HD, D₂) molecules is a reaction route that should be taken into account. Both H_2 and H abundances as well as their respective reaction efficiencies with carbon atoms determine the relative impact of both mechanisms, for which dedicated modeling will be needed. The finding that a $C + H_2$ route also leads to methane formation has the following implications.

1. The formation of CH_4 can take place at higher temperatures, e.g., 20 K instead of 10 K, because of the stronger binding of H_2 molecules to the ice surface,
2. CH_4 can be formed in the ice bulk through the interaction of entrapped H_2 with CH_n radicals obtained by dissociation of hydrocarbons caused by UV photons or cosmic-ray particles,
3. Deuterium fractionation of methane is not only dictated by D/H ratios but also by (a) the respective abundances of D₂ and HD with respect to H_2 on the surface and (b) the higher abstraction rate constant of H atoms from H_2/HD compared to D abstraction from HD/D₂; see for instance Figure 5,
4. While Qasim et al. (2020a) focused on confirming the atomic hydrogenation route of carbon to form CH_4 , here we show that not only reactions with H, but also H_2 chemistry overall, should be fully incorporated into astrochemical models. This is particularly true for models that include microscopic detail, despite the increase in computational cost.

We emphasize that astrochemical models are needed to study, in detail, the role of H_2 , HD, and D₂, for the formation of methane isotopologues under dense cloud conditions.

T.L. is grateful for support from NWO via a VENI fellowship (722.017.008). G.F. acknowledges financial support from the Russian Ministry of Science and Higher Education via the State Assignment Contract FEUZ-2020-0038. This research benefited from the financial support from the Dutch Astrochemistry Network II (DANII). Further support includes a VICI grant of NWO (the Netherlands Organization for Scientific Research). Funding by NOVA (the Netherlands Research School for Astronomy) is acknowledged.

Software: Matplotlib (Hunter 2007), Numpy (van der Walt et al. 2011), Jupyter (Kluyver et al. 2016), Chemshell (Metz et al. 2014), Molpro (Werner et al. 2012).

Appendix A Reaction Routes for the Formation of Methane Isotopologues

Below we explain the reaction pathways that lead to the formation of methane isotopologues of the form CH_nD_{4-n} with $n = 0-4$ in the experimental series 1–4. Please note that the reactions considered barrierless are reactions (1)–(4) and (6) and these are expected to take place without an isotope effect. Reaction (5) is likely possible both with H_2 and D₂, although it is currently unclear whether the rate constant is determined by the change needed in electron configuration and/or a barrier on a water-rich surface. It is possible that the reaction is slower with D₂. Reactions (10) and (11) can only take place via tunneling and, given the high barrier, these reactions are expected to be very slow with D₂. In Figure 5 three networks are depicted, analogously to Figure 4, one for each deuteration experiment. As in Figure 4, three types of reactions are considered: H/D atom addition, H_2/D_2 insertion, and H_2 abstraction. Note that we deliberately choose not to include D₂ abstraction reactions, because of the high barrier. Finally, we assume that the H or D atoms formed in situ do not take part in subsequent reactions, but desorb instead. We base this on the argument of conservation of energy and momentum (Koning et al. 2013).

The experiments are discussed below in order of increasing complexity.

A.1. Experiments 3 and 4: $C + H_2$ and/or D_2

Experiment 4A with $C + H_2$ leads clearly to the formation of CH_4 , while experiment 3A with $C + D_2$ does not lead to a CD_4 detection. At the same time experiment 3B with $C + H_2 + D_2$ shows the formation of CH_2D_2 and CH_4 , but not CD_4 . This can be understood by considering the reactions presented in the top panel of Figure 5.

A.2. Experiment 1: $C + H + H_2 + D_2$

Throughout the series for experiment 1 the only methane isotopologue species observed are CH_4 and CH_2D_2 . The reaction network in Figure 5 rationalizes these two detections by showing that these species can be formed via routes in which the majority of the reactions are barrierless. The formation of CHD_3 or CH_3D (not depicted) would need to proceed through a +D₂ tunneling abstraction reaction, which is unlikely to take place on laboratory timescales. Whether D₂ abstraction on interstellar timescales depends on the competition with diffusion needs an astrochemical model for verification.

A.3. Experiment 2: $C + H_2 + D + D_2$

In experimental series 2 we confirm the detection of CD_4 , CH_2D_2 , and a tentative detection of CH_4 . However, as can be seen from the reaction network in Figure 5, both CHD_3 and CH_3D could be formed in principle. We attribute the lack of detection of these species to:

1. the existence of a fully barrierless formation pathway for both CD_4 and CH_2D_2 ,
2. the large band strength for CH_4 , allowing a detection of small amounts,

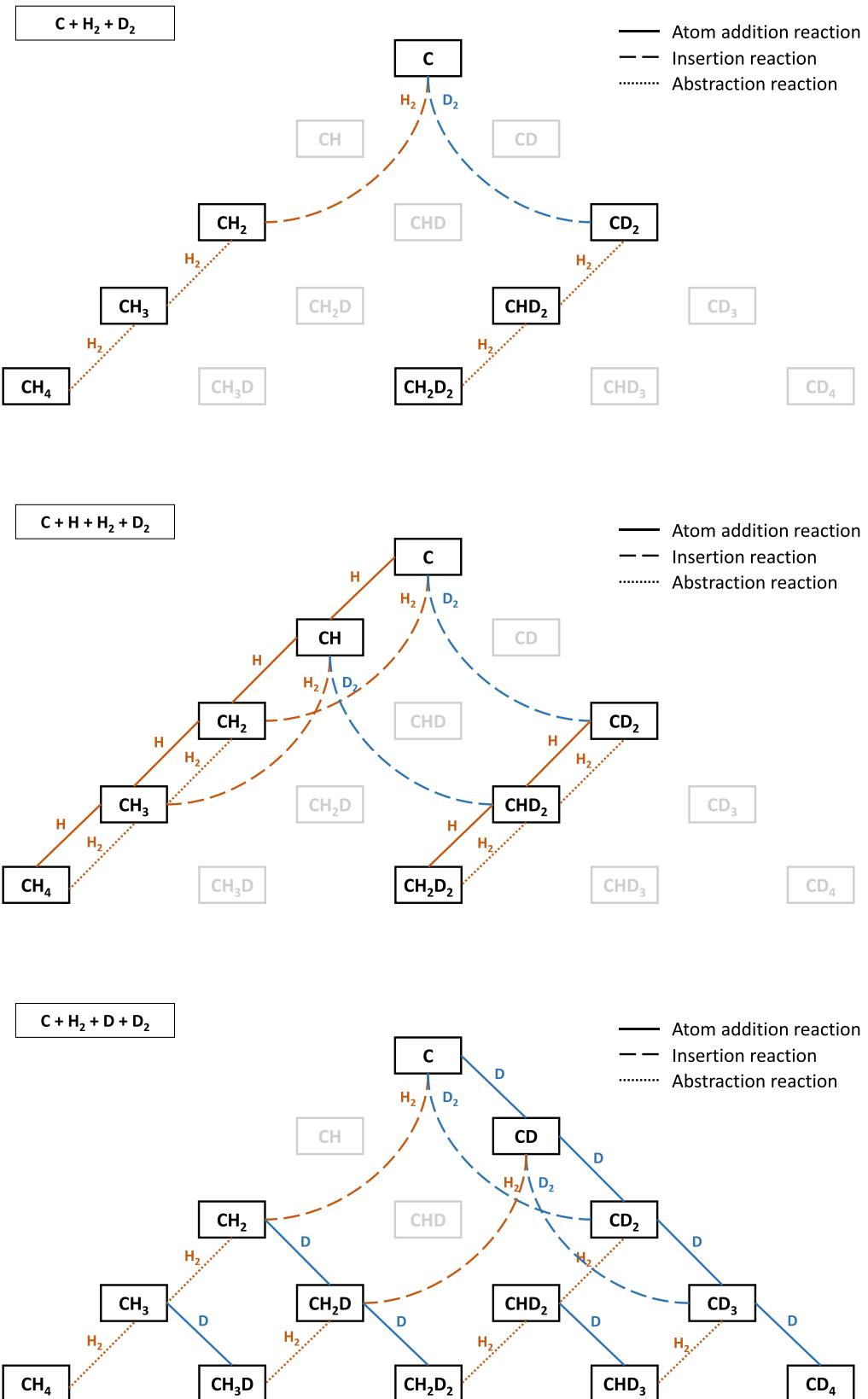


Figure 5. Reaction network for the hydrogenation and deuteration reactions leading to the formation of methane isotopologues. Analogously to Figure 4, three types of reactions are considered: H or D addition, H₂ or D₂ insertion, and H₂ abstraction.

3. the requirement for CHD_3 and CH_3D of at least one reaction that involves tunneling of the type $\text{CH}_n\text{D}_m + \text{H}_2$, which is in direct competition with a barrierless reaction of the type $\text{CH}_n\text{D}_m + \text{D}$.

For example, CH_3D can be formed via three routes, all of which contain one reaction with a barrier:

| | | | |
|----|---|---|---|
| 1. | $\text{C} + \text{H}_2 \rightarrow \text{CH}_2$ | $\text{CH}_2 + \text{H}_2 \rightarrow \text{CH}_3 + \text{H}$ | $\text{CH}_3 + \text{D} \rightarrow \text{CH}_3\text{D}$ |
| 2. | $\text{C} + \text{H}_2 \rightarrow \text{CH}_2$ | $\text{CH}_2 + \text{D} \rightarrow \text{CH}_2\text{D}$ | $\text{CH}_2\text{D} + \text{H}_2 \rightarrow \text{CH}_3\text{D} + \text{H}$ |
| 3. | $\text{C} + \text{D} \rightarrow \text{CD}$ | $\text{CD} + \text{H}_2 \rightarrow \text{CH}_2\text{D}$ | $\text{CH}_2\text{D} + \text{H}_2 \rightarrow \text{CH}_3\text{D} + \text{H}$ |

In contrast, CH_2D_2 can be formed via four routes, of which route 3 is barrierless:

| | | | |
|----|---|--|--|
| 1. | $\text{C} + \text{H}_2 \rightarrow \text{CH}_2$ | $\text{CH}_2 + \text{D} \rightarrow \text{CH}_2\text{D}$ | $\text{CH}_2\text{D} + \text{D} \rightarrow \text{CH}_2\text{D}_2$ |
| 2. | $\text{C} + \text{D}_2 \rightarrow \text{CD}_2$ | $\text{CD}_2 + \text{H}_2 \rightarrow \text{CHD}_2 + \text{H}$ | $\text{CHD}_2 + \text{H}_2 \rightarrow \text{CH}_2\text{D}_2 + \text{H}$ |
| 3. | $\text{C} + \text{D} \rightarrow \text{CD}$ | $\text{CD} + \text{H}_2 \rightarrow \text{CH}_2\text{D}$ | $\text{CH}_2\text{D} + \text{D} \rightarrow \text{CH}_2\text{D}_2$ |
| 4. | $\text{C} + \text{D} \rightarrow \text{CD}$ | $\text{CD} + \text{D} \rightarrow \text{CD}_2$ | $\text{CD}_2 + \text{H}_2 \rightarrow \text{CHD}_2 + \text{H}$ |

Appendix B Peak Positions and Detections

In Table 4 the observed peak positions are listed, along with their molecular assignment based on literature values. The last four columns indicate for which of the experiments a particular peak has been detected, with parentheses indicating a weak feature.

We confirm the results published by Qasim et al. (2020a), who showed that (deuterated) methane is efficiently formed when carbon atoms react with H (D) on a water surface via experiments 1A and 2A. Furthermore, we detect H_2CO , CO_2 , and CO in situ in the solid state. Formaldehyde is present as a product from the reaction between the carbon atom and water (Hickson et al. 2016) and its formation is the topic of another recent study (Molpeceres et al. 2021). The presence of H_2CO

further leads to the tentative detection of a CH_3OH feature in experiments 1A and 1D at 1015 cm^{-1} as a result of the hydrogenation of formaldehyde (Watanabe & Kouchi 2002; Fuchs et al. 2009; Qasim et al. 2018). Note that Fedoseev et al. (2022) showed that the reaction $\text{C} + \text{H}_2\text{CO}$ does not lead to a detectable amount of products using RAIRS, under similar experimental conditions, and therefore does not interfere with the reaction channels studied here. CO_2 is a contaminant that arises from atomic carbon sources of this design (Krasnokutski & Huisken 2014; Qasim et al. 2020b). Note also the gas-phase CO_2 bands around 2340 cm^{-1} . Qasim et al. (2020b) showed that the reaction $\text{C} + \text{CO}_2$ has a high barrier and is unlikely to take place, and Bisschop et al. (2007) have shown that the reaction $\text{H} + \text{CO}_2$ also does not take place. CO deposition as a result of the source contamination takes place with a relatively high flux

Table 4
Summary of All Detected Peak Positions, with the Exception of Water (H_2O : 3380 and 1660 cm^{-1} and D_2O : 2440 and 1220 cm^{-1})

| Peak pos. (cm^{-1}) | Molecule | Reference | Detected in Exp. 1 | Detected in Exp. 2 | Detected in Exp. 3 | Detected in Exp. 4 |
|-----------------------------------|-------------------------|-----------|-----------------------|-----------------------|-----------------------|-----------------------|
| 3007 | CH_4 | [2] | 1A, 1B | ... | ... | (4A) |
| 3000 | CH_2D_2 | [3] | (1B), 1C | ... | ... | ... |
| 2343 | CO_2 | [4] | All | All | 3A, 3B | ^a |
| 2277 | CH_2D_2 | [3] | 1C | (2C) | ... | ... |
| 2250 | CD_4 | [5] | ... | 2A, 2B | ... | ... |
| 2226 | CH_2D_2 | [3] | 1C | ... | ... | ... |
| 2152 | CO | [6] | All | All | 3A, 3B | All |
| 2137 | CO | [6] | All | All | 3A, 3B | All |
| 1717 | H_2CO | [7] | All | All | 3A, 3B | ... |
| 1666 | D_2CO | [8] | ... | ... | ... | 4A, 4B |
| 1500 | H_2CO | [7] | All | All | 3A, 3B | ... |
| 1430 | CH_2D_2 | [3] | 1B, 1C | ... | ... | ... |
| 1303 | CH_4 | [1, 2] | 1A, 1B, (1C) | (2C) | (3B) | 4A |
| 1250 | H_2CO | [7] | 1A | ... | 3A, 3B | ... |
| 1231 | CH_2D_2 | [3] | 1B, (1B) | ... | ... | ... |
| 1102 | D_2CO | [8] | ... | ... | ... | (4A), 4B |
| 1083 | CH_2D_2 | [3] | 1B, | (2C) | 3B | ... |
| 1028 | CH_2D_2 | [3] | 1B, (1B) | (2A), 2B, 2C | 3B | ... |
| 1015 | CH_3OH | [9] | 1A, (1B) | ... | ... | ... |
| 993 | CD_4 | [5] | (1C) | 2A, 2B, (2C) | ... | ... |
| 991 | D_2CO | [8] | ... | ... | ... | 4A, 4B |

Notes. Experiments in parentheses indicate a weak feature or tentative detection.

^a Overlaps with the D_2O stretch mode.

References. [1] Shimanouchi (1972); [2] Hagen et al. (1983); Quattroci & Ewing (1992a); [3] Quattroci & Ewing (1992b); [4] Gerakines et al. (1995); [5] Chapados & Cabana (1972); Edling et al. (1987); [6] Schmitt et al. (1989); [7] Schutte et al. (1996); [8] Tso & Lee (1984); Nagaoka et al. (2005); [9] Qasim et al. (2018).

of $8 \times 10^{11} \text{ cm}^{-2} \text{ s}^{-1}$, and subsequent reactions with H and C can lead to the formation of ketene (Fedoseev et al. 2022). This does not, however, interfere with the formation routes of methane since the reaction channels are separated. Finally, reactions between H₂O and CH_x radicals are not expected to play a role, because of the high barriers or endothermicity of the reaction involved (Jursic 1998; Tzeli & Mavridis 2005; Bergeat et al. 2009; Lamberts et al. 2017; Qasim et al. 2020a).

Appendix C Potential Energy Surfaces for the C_{2v} and C_s Symmetries

Figure 6 shows the potential energy cuts for the reaction C + H₂ → CH₂, reaction (5), on both C_{2v} and C_s surfaces. Note that for both C_s surfaces at R > 1.8 Å we faced convergence issues. These values have been omitted. The position of the ³CH₂ ground state is indicated in each figure.

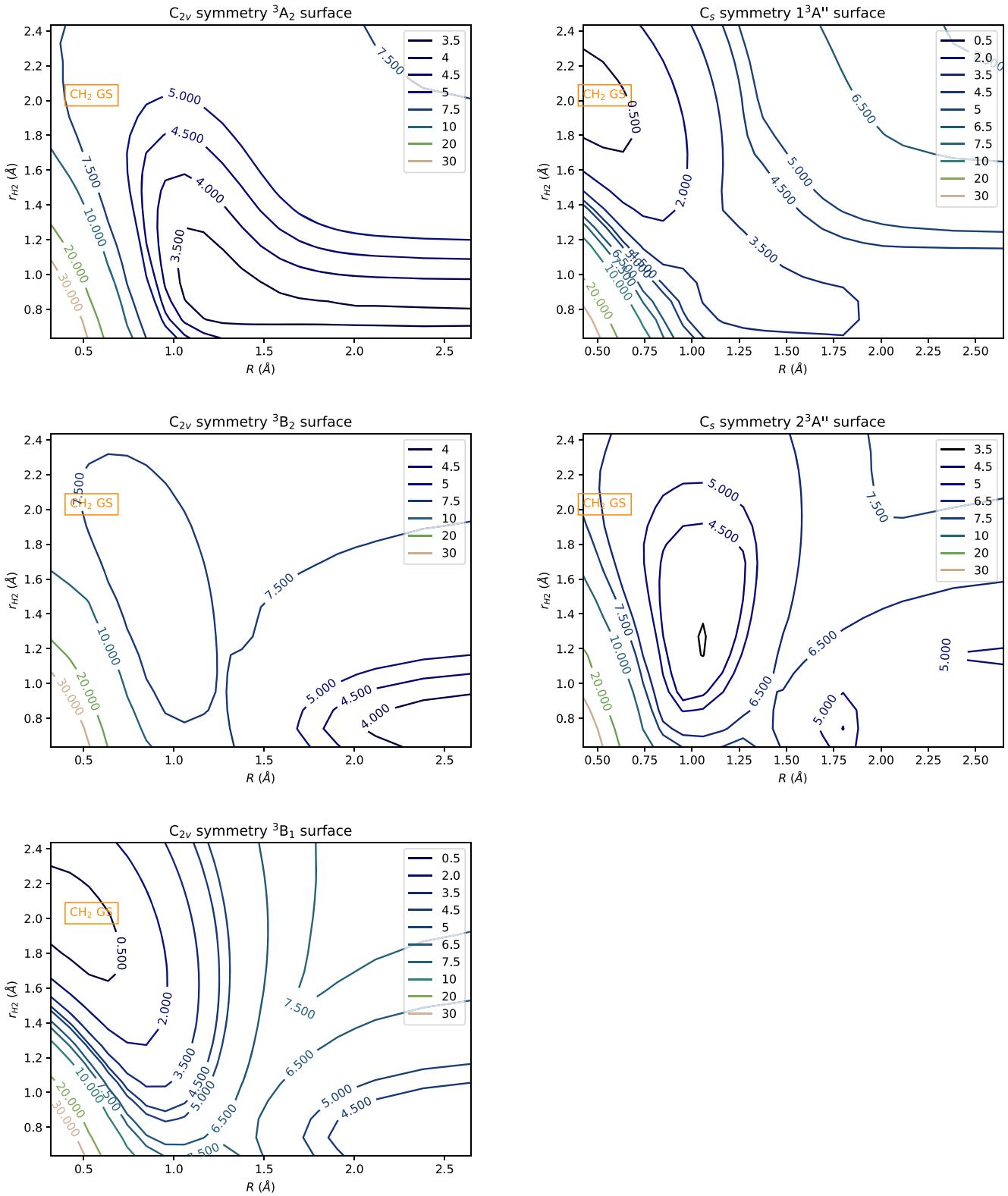


Figure 6. Potential energy cuts for the reaction $C + H_2$ on three C_{2v} (left column) and two C_s symmetry (right column) surfaces. Contour units are in eV.

ORCID iDs

- Thanja Lamberts <https://orcid.org/0000-0001-6705-2022>
 Gleb Fedoseev <https://orcid.org/0000-0003-2434-2219>
 Danna Qasim <https://orcid.org/0000-0002-3276-4780>
 Ko-Ju Chuang <https://orcid.org/0000-0001-6877-5046>
 Julia C. Santos <https://orcid.org/0000-0002-3401-5660>
 Harold Linnartz <https://orcid.org/0000-0002-8322-3538>

References

- Adler, T. B., Knizia, G., & Werner, H.-J. 2007, *JChPh*, **127**, 221106
 Albar, J. D., Summerfield, A., Cheng, T. S., et al. 2017, *NatSR*, **7**, 6598
 Anton, R., Wiegner, T., Naumann, W., et al. 2000, *RSci*, **71**, 1177
 Baskin, C. P., Bender, C. F., Bauschlicher, C. W., & Schaefer, H. F. 1974, *JChS*, **96**, 2709
 Bauschlicher, C. W. 1978, *CPL*, **56**, 31
 Bauschlicher, C. W., Haber, K., Schaefer, H. F., & Bender, C. F. 1977, *JChS*, **99**, 3610
 Becke, A. D. 1988, *PhRvA*, **38**, 3098
 Becke, A. D. 1993, *JChPh*, **98**, 5648
 Bergeat, A., Moisan, S., Méreau, R., & Loison, J.-C. 2009, *CPL*, **480**, 21
 Beyer, A. N., Richardson, J. O., Knowles, P. J., Rommel, J., & Althorpe, S. C. 2016, *J. Phys. Chem. Lett.*, **7**, 4374
 Bisbas, T. G., Schruba, A., & van Dishoeck, E. F. 2019, *MNRAS*, **485**, 3097
 Bisschop, S. E., Fuchs, G. W., van Dishoeck, E. F., & Linnartz, H. 2007, *A&A*, **474**, 1061
 Boogert, A. A., Gerakines, P. A., & Whittet, D. C. 2015, *ARA&A*, **53**, 541
 Boogert, A. C. A., Schutte, W. A., Tielens, A. G. G. M., et al. 1996, *A&A*, **315**, L377
 Brown, P. D., & Charnley, S. B. 1991, *MNRAS*, **249**, 69
 Brown, P. D., Charnley, S. B., & Millar, T. J. 1988, *MNRAS*, **231**, 409
 Burton, M. G., Ashley, M. C. B., Braidling, C., et al. 2015, *ApJ*, **811**, 13
 Callan, C. G., & Coleman, S. 1977, *PhRvD*, **16**, 1762
 Chapados, C., & Cabana, A. 1972, *CajCh*, **50**, 3521
 Chuang, K. J., Fedoseev, G., Qasim, D., et al. 2018, *A&A*, **617**, A87
 Cuppen, H. M., & Herbst, E. 2007, *ApJ*, **668**, 294
 D'Hendecourt, L. B., Allamandola, L. J., & Greenberg, J. M. 1985, *A&A*, **152**, 130
 Duchovic, R. J., & Hase, W. L. 1985, *JChPh*, **82**, 3599
 Dufort, D., Toubin, C., & Monnerville, M. 2021, *FrASS*, **8**, 24
 Dunning, T. H., Peterson, K. A., & Wilson, A. K. 2001, *JChPh*, **114**, 9244
 Dunning, T. H. J. 1989, *JChPh*, **90**, 1007
 Edling, J. A., Richardson, H. H., & Ewing, G. E. 1987, *JMoSt*, **157**, 167
 Fedoseev, G., Qasim, D., Chuang, K.-J., et al. 2022, *ApJ*, **924**, 110
 Fredon, A., Groenenboom, G. C., & Cuppen, H. M. 2021, *ESC*, **5**, 2032
 Fuchs, G. W., Cuppen, H. M., Ioppolo, S., et al. 2009, *A&A*, **505**, 629
 Furuya, K., Aikawa, Y., Hincelin, U., et al. 2015, *A&A*, **584**, A124
 Gamallo, P., Defazio, P., Akpinar, S., & Petrongolo, C. 2012, *JPCA*, **116**, 8291
 Garrod, R. T. 2013, *ApJ*, **778**, 158
 Gerakines, P. A., Schutte, W. A., Greenberg, J. M., & van Dishoeck, E. F. 1995, *A&A*, **296**, 810
 Goldsmith, P. F., & Li, D. 2005, *ApJ*, **622**, 938
 González, M., Saracibar, A., & García, E. 2011, *PCCP*, **13**, 3421
 Greenler, R. G. 1966, *JChPh*, **44**, 310
 Hagen, W., Tielens, A. G. G. M., & Greenberg, J. M. 1983, *A&AS*, **51**, 389
 Harding, L., Guadagnini, R., & Schatz, G. 1993, *JPhCh*, **97**, 5472
 Hasegawa, T. I., & Herbst, E. 1993, *MNRAS*, **261**, 83
 Henning, T. K., & Krasnokutski, S. A. 2019, *NatAs*, **3**, 568
 Hickson, K. M., Loison, J.-C., Lique, F., & Klos, J. 2016, *JPCA*, **120**, 2504
 Hickson, K. M., Loison, J.-C., Nuñez-Reyes, D., & Méreau, R. 2016, *J. Phys. Chem. Lett.*, **7**, 3641
 Hiraoka, K., Miyagoshi, T., Takayama, T., Yamamoto, K., & Kihara, Y. 1998, *ApJ*, **498**, 710
 Hunter, J. D. 2007, *CSE*, **9**, 90
 Ioppolo, S., Cuppen, H. M., Romanzin, C., van Dishoeck, E. F., & Linnartz, H. 2008, *ApJ*, **686**, 1474
 Ioppolo, S., Fedoseev, G., Lamberts, T., Romanzin, C., & Linnartz, H. 2013, *RSci*, **84**, 073112
 Janssen, C. L., & Nielsen, I. M. 1998, *CPL*, **290**, 423
 Jursic, B. S. 1998, *JPCA*, **102**, 9255
 Kästner, J., Carr, J. M., Keal, T. W., et al. 2009, *JPCA*, **113**, 11856
 Keene, J., Blake, G. A., Phillips, T. G., Huggins, P. J., & Beichman, C. A. 1985, *ApJ*, **299**, 967
 Kendall, R. A., Dunning, T. H., & Harrison, R. J. 1992, *JChPh*, **96**, 6796
 Kim, G.-S., Nguyen, T. L., Mebel, A. M., Lin, S. H., & Nguyen, M. T. 2003, *JPCA*, **107**, 1788
 Kluyver, T., Ragan-Kelley, B., Pérez, F., et al. 2016, in Positioning and Power in Academic Publishing: Players, Agents and Agendas, ed. F. Loizides & B. Schmidt (Amsterdam: IOS Press), 87
 Knizia, G., Adler, T. B., & Werner, H.-J. 2009, *JChPh*, **130**, 054104
 Koning, J., Kroes, G. J., & Arasa, C. 2013, *JChPh*, **138**, 104701
 Krasnokutski, S. A., & Huisken, F. 2014, *ApPhL*, **105**, 113506
 Krasnokutski, S. A., Kuhn, M., Renzler, M., et al. 2016, *ApJL*, **818**, L31
 Lacy, J. H., Carr, J. S., Evans, N. J. I., et al. 1991, *ApJ*, **376**, 556
 Lambert, N., Kaltsoyannis, N., Price, S. D., Žabka, J., & Herman, Z. 2006, *JPCA*, **110**, 2898
 Lamberts, T., Cuppen, H. M., Ioppolo, S., & Linnartz, H. 2013, *PCCP*, **15**, 8287
 Lamberts, T., Fedoseev, G., Kästner, J., Ioppolo, S., & Linnartz, H. 2017, *A&A*, **599**, A132
 Lamberts, T., Samanta, P. K., Köhn, A., & Kästner, J. 2016, *PCCP*, **18**, 33021
 Langer, W. 1976, *ApJ*, **206**, 699
 Lee, C., Yang, W., & Parr, R. G. 1988, *PhRvB*, **37**, 785
 Li, J., Chen, J., Zhao, Z., et al. 2015, *JChPh*, **142**, 204302
 McIlroy, A., & Tully, F. P. 1993, *JChPh*, **99**, 3597
 Medvedev, D. M., Harding, L. B., & Gray, S. K. 2006, *MolPh*, **104**, 73
 Meisner, J., Lamberts, T., & Kästner, J. 2017, *ESC*, **1**, 399
 Metz, S., Kästner, J., Sokol, A. A., Keal, T. W., & Sherwood, P. 2014, *Wiley Interdiscip. Rev. Comput. Mol. Sci.*, **4**, 101
 Miller, K. J. 1975, *JChPh*, **62**, 1759
 Miyauchi, N., Hidaka, H., Chigai, T., et al. 2008, *CPL*, **456**, 27
 Molpeceres, G., Kästner, J., Fedoseev, G., et al. 2021, *J. Phys. Chem. Lett.*, **12**, 10854
 Molpeceres, G., Zaverkin, V., & Kästner, J. 2020, *MNRAS*, **499**, 1373
 Murrell, J. N., Pedley, J. B., & Durmaz, S. 1973, *J. Chem. Soc., Faraday Trans.*, **2**, 1370
 Nagaoka, A., Watanabe, N., & Kouchi, A. 2005, *ApJL*, **624**, L29
 Öberg, K. I., Boogert, A. C. A., Pontoppidan, K. M., et al. 2008, *ApJ*, **678**, 1032
 Papadopoulos, P. P., Thi, W. F., & Viti, S. 2004, *MNRAS*, **351**, 147
 Peterson, K. A., Adler, T. B., & Werner, H.-J. 2008, *JChPh*, **128**, 084102
 Qasim, D., Chuang, K.-J., Fedoseev, G., et al. 2018, *A&A*, **612**, A83
 Qasim, D., Fedoseev, G., Chuang, K. J., et al. 2020a, *NatAs*, **4**, 781
 Qasim, D., Witlox, M. J. A., Fedoseev, G., et al. 2020b, *RSci*, **91**, 054501
 Quattroci, L. M., & Ewing, G. E. 1992a, *JChPh*, **96**, 4205
 Quattroci, L. M., & Ewing, G. E. 1992b, *CPL*, **197**, 308
 Richardson, J. O. 2016, *JChPh*, **144**, 114106
 Rommel, J. B., & Kästner, J. 2011, *JChPh*, **134**, 184107
 Schmidt, A., Offermann, J., & Anton, R. 1996, *TSF*, **281**, 105
 Schmitt, B., Greenberg, J. M., & Grim, R. J. A. 1989, *ApJL*, **340**, L33
 Schutte, W. A., Gerakines, P. A., Geballe, T. R., van Dishoeck, E. F., & Greenberg, J. M. 1996, *A&A*, **309**, 633
 Shimanouchi, T. 1972, *Nat. Stand. Ref. Data Ser. Nat. Bur. Stand.*, **1**, 39
 Shimonishi, T., Nakatani, N., Furuya, K., & Hama, T. 2018, *ApJ*, **855**, 27
 Simončič, M., Semenov, D., Krasnokutski, S., Henning, T., & Jäger, C. 2020, *A&A*, **637**, A72
 Snow, T. P., & McCall, B. J. 2006, *ARA&A*, **44**, 367
 Tso, T. L., & Lee, E. K. C. 1984, *JPhCh*, **88**, 5475
 Tzeli, D., & Mavridis, A. 2005, *IQC*, **104**, 497
 van de Hulst, H. C. 1946, *RAOU*, **11**, 2
 van de Hulst, H. C. 1949, The Solid Particles in Interstellar Space (Utrecht: Drukkerij Schotanus & Jens)
 van der Walt, S., Colbert, S. C., & Varoquaux, G. 2011, *CSE*, **13**, 22
 van Dishoeck, E. F., & Black, J. H. 1988, *ApJ*, **334**, 771
 van Harreveld, R., van Hemert, M. C., & Schatz, G. C. 2002, *JChPh*, **116**, 6002
 Vasyunin, A. I., & Herbst, E. 2013, *ApJ*, **762**, 86
 Wakelam, V., Loison, J. C., Herbst, E., et al. 2015, *ApJS*, **217**, 20
 Wakelam, V., Loison, J. C., Mereau, R., & Ruaud, M. 2017, *MolAs*, **6**, 22
 Watanabe, N., & Kouchi, A. 2002, *ApJL*, **571**, L173
 Weigend, F., & Ahlrichs, R. 2005, *PCCP*, **7**, 3297
 Werner, H., & Knowles, P. J. 1988, *JChPh*, **89**, 5803
 Werner, H.-J., Knowles, P. J., Knizia, G., et al. 2018, MOLPRO, v2018.1, a package of ab initio programs, <https://www.molpro.net/>
 Werner, H.-J., Knowles, P. J., Knizia, G., Manby, F. R., & Schütz, M. 2012, *WIREs Comput. Mol. Sci.*, **2**, 242
 Werner, H.-J., Knowles, P. J., Manby, F. R., et al. 2020, *JChPh*, **152**, 144107
 Woon, D. E., & Dunning, T. H. 1993, *JChPh*, **98**, 1358