

Surface formation of CO₂ ice at low temperatures

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ABSTRACT

The surface formation of CO₂ at low temperatures through the reaction CO + OH and direct dissociation of the resulting HO–CO complex is shown by hydrogenation of a CO:O₂ ice mixture. Such a binary ice is not fully representative for an interstellar ice, but the hydrogenation of O₂ ice produces OH radicals, which allows the investigation of the interstellar relevant CO + OH solid state reaction under fully controlled laboratory conditions. Similar recent astrophysical ice studies have focused on the investigation of isolated surface reaction schemes, starting from the hydrogenation of pure ices, like solid CO or O₂. For such ices, no CO₂ formation is observed upon H-atom exposure. The hydrogenation of binary ice mixtures presented here allows to investigate for the first time the influence of the presence of other species in the ice on the pure ice reaction schemes. Mixtures of CO:O₂ are deposited on a substrate in an ultra high vacuum setup at low temperatures (15 and 20 K) and subsequently hydrogenated. The ice is monitored by means of Reflection Absorption InfraRed Spectroscopy (RAIRS). Results show that solid CO₂ is formed in all studied CO:O₂ mixtures under our laboratory conditions. Within the experimental uncertainties no dependency on ice temperature or composition is observed. The laboratory results show a correlation between the formation of CO₂ and H₂O, which is consistent with the astronomical observation of solid CO₂ in water-rich environments. The results also show that the contemporary presence of CO and O₂ molecules in the ice influences the final product yields of the separate CO + H (H₂CO, CH₃OH) and O₂ + H (H₂O₂ and H₂O) channels, even though the formation rates are not significantly affected.

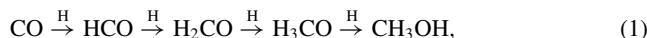
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1 INTRODUCTION

Infrared Space Observatory and Spitzer Space Telescope observations have shown that H₂O, CO, CO₂ and, in some cases, CH₃OH represent the bulk of solid-state species in dense molecular clouds and star-forming regions. Other ice components, such as CH₄, NH₃, OCN[−], H₂CO, HCOOH, SO₂ and OCS have abundances <5 per cent relative to H₂O (e.g. Gibb et al. 2004; Boogert et al. 2008; Pontoppidan et al. 2008; Öberg et al. 2008; Zasowski et al. 2009; Bottinelli et al. 2010). Several of these species are assumed to be formed in solid state reactions on the surfaces of icy dust grains, as outlined by Tielens & Hagen (1982). Although these reactions have been postulated nearly 30 yr ago, few have been measured in the laboratory at low temperatures and under ultra high vacuum (UHV)

conditions until recently. Over the past decade, detailed laboratory experiments have started to investigate isolated surface reaction schemes, starting from the hydrogenation of simple and pure ices, like solid CO or O₂.

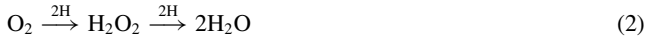
Several groups proved that the hydrogenation of CO ice at low temperatures (12–20 K) leads to the subsequent formation of H₂CO and CH₃OH (e.g. Watanabe et al. 2004, 2006; Fuchs et al. 2009). The experiments showed that this hydrogenation process involves only the upper monolayers (~4 ML, where 1 ML corresponds to 10¹⁵ molecules cm^{−2}) of the CO ice and formation rates drop at temperatures higher than 15 K, since the desorption of H atoms becomes important at these temperatures. The hydrogenation of CO to CH₃OH proceeds in four steps,



where the first step from CO to HCO and the third step from H₂CO to H₃CO have a barrier.

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The other surface reaction channel that has been well investigated is the hydrogenation of O₂ ice, which leads to the formation of H₂O₂ and H₂O (e.g. Ioppolo et al. 2008, 2010; Miyauchi et al. 2008; Cuppen et al. 2010). This hydrogenation process



behaves differently compared to the hydrogenation of CO ice (reaction scheme 2 shows the simplified version of this route, as discussed by Tielens & Hagen 1982). In this case, the penetration depth of H atoms in the O₂ ice increases with temperature, even at values close to the desorption temperature of the O₂ layer, involving the bulk of the ice (tens of monolayers). Thus, H atoms trapped in the ice can diffuse and eventually react up to much higher temperatures. Moreover, at least the formation of H₂O₂ does not exhibit any noticeable barrier.

The present work is a further step towards a laboratory investigation of interstellar relevant surface reactions in more complex ices by studying for the first time the formation of solid CO₂ through hydrogenation of a binary CO:O₂ ice mixture as well as investigating the competition between the two separate hydrogenation channels (CO + H and O₂ + H).

CO₂ is one of the most common and abundant ices (d’Hendecourt & Jourdain de Muizon 1989), yet its formation routes are still uncertain. It is widely accepted that CO₂ is not formed efficiently in the gas phase, with subsequent accretion on to the grains ($\text{CO}_2^{\text{gas}}/\text{CO}_2^{\text{ice}} \ll 1$; van et al. 1996; Boonman et al. 2003). Therefore, the observed CO₂ most likely has to be formed in the solid phase. Several reaction mechanisms have been proposed with an efficiency depending on astronomical environment. Energetic processing, such as UV and ion irradiation of interstellar ice analogues, has been investigated in various laboratories and proposed as an efficient CO₂ formation mechanism (e.g. Hagen, Allamandola & Greenberg 1979; Mennella, Palumbo & Baratta 2004; Loeffler et al. 2005; Mennella et al. 2006; Ioppolo et al. 2009). Furthermore, in the absence of UV irradiation, several cold solid-phase reaction channels have been reported in the past decades as an alternative formation mechanism to explain the CO₂ abundance observed in cold clouds (e.g. Tielens & Hagen 1982; d’Hendecourt et al. 1985; Grim & d’Hendecourt 1986; Ruffle & Herbst 2001; Fraser & van Dishoeck 2004; Stantcheva & Herbst 2004; Goumans, Uppal & Brown 2008; Goumans & Andersson 2010). The most straightforward surface reaction channel is the addition of an O atom to solid CO ice. The reaction CO + O → CO₂ has been experimentally investigated only by temperature programmed desorption experiments using thermal O atoms below 160 K (Roser et al. 2001) and by energetic O atoms (Madzunkov et al. 2006). This surface reaction channel has a high reaction barrier (Grim & d’Hendecourt 1986), because the CO(¹Σ) + O(³P) reactants do not correlate directly with the singlet ground state CO₂(¹Σ) (2970 K in the gas phase; Talbi, Chandler & Rohl 2006). Ruffle & Herbst (2001) found in their astrochemical model that they were only able to reproduce the CO₂ abundances observed towards the cold (10 K) cloud Elias 16, if they artificially lowered the barrier to 130 K. Recently, Goumans & Andersson (2010) used harmonic quantum transition state theory to conclude that whilst quantum mechanical tunnelling through the activation barrier increases the classical reaction rate for reaction CO + O at low temperatures (10–20 K), the onset of tunnelling is at too low temperatures for the reaction to efficiently contribute to CO₂ formation in quiescent clouds.

Solid CO₂ is further suggested to be formed through the surface reaction HCO + O, which presents two exit channels (CO₂ + H and CO + OH; Ruffle & Herbst 2001). Alternatively,

solid CO₂ can be formed through the surface reaction CO + OH, which yields a HO–CO intermediate. This complex can directly dissociate, forming solid CO₂ and leaving a H atom, or can be stabilized by intramolecular energy transfer to the ice surface and eventually react with an incoming H atom in a barrierless manner to form CO₂ and H₂ (Goumans et al. 2008).

Fig. 1 shows a schematic representation of all the reaction networks investigated in this work (solid arrows) and links the previously studied CO + H and O₂ + H channels through the observed CO₂ formation. The dashed and dotted arrows represent suggested CO₂ formation routes in the networks of Tielens & Hagen (1982) and Ruffle & Herbst (2001), that are not experimentally confirmed at low temperature in our studies.

In Section 4.1 we discuss that CO₂ is formed under our experimental conditions through the reaction CO + OH. Here OH radicals are formed through the hydrogenation of O₂ ice, while in space they can also result from the O + H reaction or from the photodissociation of H₂O ice. Recently, Oba et al. (2010) also investigated the reaction CO + OH depositing CO molecules on a cold substrate (10 and 20 K) together with H₂O fragments (OH, H, O and H₂) produced by dissociating H₂O molecules in a microwave source. They observed the formation of solid CO₂ at low temperature by using CO and OH beams to initiate surface reactions on a cold substrate. Their experiments differ from ours. In the present work, we hydrogenate CO:O₂ ices and OH radicals are produced in the ice through the O₂ + H channel, providing additional information on the interaction between different surface reaction schemes. In a previous paper (Ioppolo et al. 2011), we investigated experimentally the hydrogenation of the HO–CO complex, which has three exit channels (CO₂ + H₂, HCOOH, H₂O + CO) with a purely statistical branching ratio as suggested by density functional theory models and in agreement with our experimental results (Goumans et al. 2008; Ioppolo et al. 2011). In the present study, the HO–CO complex itself is not observed since it is efficiently dissociated under our experimental conditions to form CO₂. More details are reported in Section 4. In Sections 2 and 3, the experimental method and data analysis are discussed.

2 EXPERIMENTAL DETAILS

The experiments are performed using an UHV setup, which consists of a main chamber (10^{−10} mbar) and an atomic beam line. Details are available from Fuchs et al. (2009) and Ioppolo et al. (2010). The ice is grown on a gold coated copper substrate (12–300 K) that is mounted on the cold head of a close-cycle He cryostat. Deposition of selected ¹²C¹⁶O:¹⁶O₂ mixtures (4:1, 1:1 and 1:4) proceeds under an angle of 45° and with a rate of 0.7 ML min^{−1}. After deposition the ice mixture is exposed to a cold H-atom beam. H₂ molecules are dissociated into the capillary of a well-characterized thermal cracking source (Tschersich & von Bonin 1998; Tschersich 2000; Tschersich, Fleischhauer & Schuler 2008), which is used to hydrogenate the sample. A quartz pipe with a nose-shaped form is placed along the path of the dissociated beam to efficiently thermalize all H atoms to room temperature through surface collisions before they reach the ice sample. In this way, hot species (H; H₂) cannot reach the ice directly. Furthermore, the relatively high temperature of 300 K of the incident H atoms in our experiments does not affect the experimental results, since H atoms are thermally adjusted to the surface temperature as has been shown in Fuchs et al. (2009). The final H-atom flux

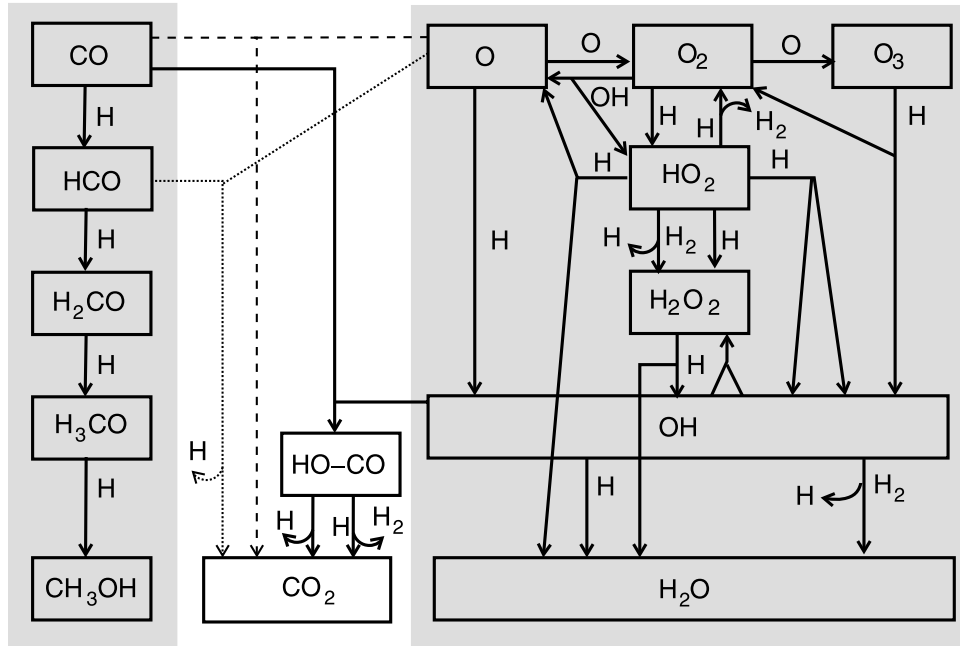


Figure 1. A schematic representation of the reaction network as discussed in the present study. The CO + H channel is shown on the left-hand side of the figure, while the O/O₂/O₃ + H channels are plotted as presented in Cuppen et al. (2010) on the right-hand side. The possible CO₂ formation routes are shown in between the CO + H and O/O₂/O₃ + H channels: the dissociation of the HO–CO intermediate (solid arrow) is one of the topics of this work; the hydrogenation of the HO–CO complex (solid arrow) is presented in Ioppolo et al. (2011); the suggested CO + O (dashed arrow) and HCO + O (dotted arrow) routes are not experimentally confirmed at low temperature (Tielens & Hagen 1982; Ruffe & Herbst 2001).

(2.5×10^{13} atoms cm⁻² s⁻¹) is measured at the substrate position in the main chamber using a quadrupole mass spectrometer, following the procedure as described in the appendix of Ioppolo et al. (2011). The error in the absolute H-atom flux determination is within 50 per cent. Ices are monitored by means of Reflection Absorption InfraRed Spectroscopy (RAIRS) using a Fourier Transform InfraRed spectrometer (FTIR), which covers the range between 4000 and 700 cm⁻¹ (2.5–14 μm). A spectral resolution of 1 cm⁻¹ is used and 128 scans are co-added for one spectrum. RAIR difference spectra (ΔA) relative to the initial unprocessed CO:O₂ ice are acquired every few minutes during H-atom exposure.

We performed a control experiment at 15 K in which a CO:O₂ ice is exposed to an H₂ molecular beam (i.e. without H atoms) to show that the products detected in the hydrogenation experiments are formed on the surface and do not originate from background deposition. Only small amounts of H₂O are detected in this experiment, which gives us an estimate for the background contamination, which is negligible. None of the other products is detected in this way. The present experiments use the same H-atom flux as adopted in Fuchs et al. (2009) and Ioppolo et al. (2011). Therefore, the hydrogenation of our mixtures involves effectively a lower H-atom flux per (CO + H and O₂ + H) reaction channel.

The interstellar solid CO:O₂ mixing ratio is observationally constrained to > 1:1 (e.g. Vandebussche et al. 1999; Pontoppidan et al. 2003). This conclusion was deduced earlier by d’Hendecourt et al. (1985) on the basis of a simple interacting gas-phase/solid-phase time-dependent model. In our experiments, the binary mixture ratios (CO:O₂ = 4:1, 1:1 and 1:4) are selected to efficiently produce thermal OH radicals in the ice in order to investigate the formation of CO₂. These experiments are also meant to test the interaction of the aforementioned individual surface reaction channels, rather than simulating a complete realistic interstellar ice evolution.

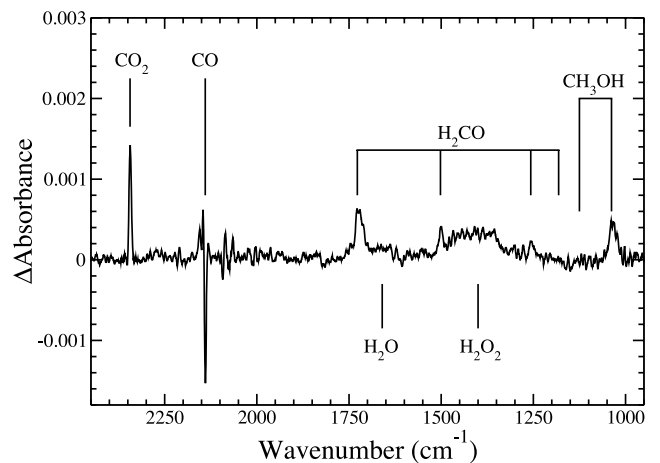


Figure 2. RAIR difference spectrum of the CO:O₂ = 1:4 ice, with respect to the spectrum before H-atom addition, at 15 K after a H-atom fluence of 1.3×10^{17} atoms cm⁻².

3 DATA ANALYSIS

Fig. 2 shows a RAIR difference spectrum of a CO:O₂ = 1:4 mixture acquired after a H-atom fluence (flux × time) of 1.3×10^{17} atoms cm⁻². The negative peak shown in Fig. 2 is caused by the CO use-up in surface reaction processes. O₂ is infrared inactive and therefore cannot be observed in this spectrum. All the final reaction products obtained from the hydrogenation of a pure CO ice (H₂CO and CH₃OH; e.g. Watanabe et al. 2004, 2006; Fuchs et al. 2009) and a pure O₂ ice (H₂O and H₂O₂; e.g. Ioppolo et al. 2008; Miyauchi et al. 2008) are present. Neither the intermediate species from the separate CO and O₂ channels, like HCO, H₃CO, HO₂, OH and O₃,

nor more complex species, like the stabilized HO–CO intermediate, HCOOH and H₂CO₃,¹ are observed as was the case in experiments by Ioppolo et al. (2011) in which the CO:O₂ mixtures and H atoms were deposited simultaneously. However, a new feature appears at $\sim 2344\text{ cm}^{-1}$, which belongs to the asymmetrical stretching mode of CO₂ ice. This molecule is formed in all our performed experiments at 15 and 20 K, with different CO:O₂ mixing ratios, but was not found in the previous CO + H or O₂ + H experiments.

The infrared spectra are reduced to obtain the column densities of the newly formed species. As a first step in the infrared data analysis, a straight baseline is subtracted from all spectra. Some absorption features, like the H₂O bending mode ($\sim 1650\text{ cm}^{-1}$) and the H₂CO $\nu(\text{CO})$ stretching mode ($\sim 1720\text{ cm}^{-1}$) suffer from spectral overlap. Here a multi-Gaussian fit is used to determine the area of the selected bands. Since the asymmetric 1440 cm^{-1} H₂O₂ band overlaps with the 1500 cm^{-1} H₂CO band, a spectrum of pure H₂O₂ ice is fitted in addition to a Gaussian. The spectrum of solid H₂O₂ is obtained as discussed by Cuppen et al. (2010), by co-depositing H atoms and O₂ molecules with a ratio of H/O₂ = 20 and subsequently heating the ice to a temperature higher than 30 K, which is just above the O₂ desorption temperature (Acharyya et al. 2007).

The column density N_X (molecules cm^{-2}) of species X in the ice is calculated using: $N_X = \int A(\nu) d\nu / S_X$, where $A(\nu)$ is the wavelength-dependent absorbance. Since literature values of transmission band strengths cannot be used in reflection measurements, an apparent absorption band strength, S_X of species X , is determined by individual calibration experiments. These have been described in detail in Fuchs et al. (2009), Ioppolo et al. (2010) and Cuppen et al. (2010). Like for CO, CH₃OH and H₂O, an isothermal desorption experiment has been performed to determine the apparent absorption band strength of CO₂ by determining the transition from zeroth-order to first-order desorption. This is assumed to occur at the onset of the submonolayer regime and appears in the desorption curve as a sudden change in slope. Since pure H₂CO and H₂O₂ are experimentally difficult to deposit, because of their chemical instability, the values for $S_{\text{H}_2\text{CO}}$ and $S_{\text{H}_2\text{O}_2}$ are obtained by assuming mass balance as reported in Fuchs et al. (2009) and Cuppen et al. (2010), respectively.

4 RESULTS AND DISCUSSION

4.1 Formation of solid CO₂

Fig. 3 shows the CO₂ column density as a function of the H-atom fluence, confirming the CO₂ formation for the three different mixing ratios and two temperatures investigated. Here neither the CO₂ formation rate nor its final yield depends significantly on either temperature or mixing ratio for the studied values. Such a behaviour is unexpected, since the separate reaction routes CO + H and O₂ + H clearly depend on temperature, as shown by Fuchs et al. (2009) and Ioppolo et al. (2010). The limiting factor for the CO₂ synthesis in our experiments is, therefore likely, the amount of ice that can be penetrated by H atoms, which is only a few monolayers. This is caused by the presence of CO molecules affecting the penetration depth of the H atoms in the ice (Fuchs et al. 2009). Thus, the amount of CO₂ formed in all our experiments is always less than a monolayer. CO₂ subsequently does not contribute to further molecular synthesis in the ice upon ongoing hydrogenation. Bisschop et al. (2007) showed

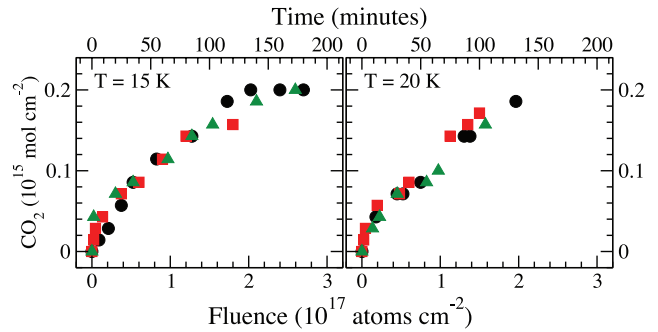


Figure 3. CO₂ column density as a function of the H-atom fluence and time of H-atom exposure at 15 K (left) and 20 K (right) for the three mixtures studied: CO:O₂ = 4:1 (circle), 1:1 (square), and 1:4 (triangle).

experimentally that CO₂ does not react with H atoms and is a stable molecule under interstellar ice analogue conditions.

Fig. 1 summarizes schematically the reaction network which leads to the formation of solid CO₂ starting from the combination of the CO + H and O₂ + H channels. Analysing the species present in our ice after H-atom addition, we can identify which reaction channel is most likely responsible for the formation of solid CO₂ ice. The hydrogenation of the HO–CO intermediate (black arrow in the centre of Fig. 1) should not occur in our experiments, since HCOOH is not detected in the infrared spectra. Density functional calculations (Goumans et al. 2008), confirmed by our previous experimental results (Ioppolo et al. 2011), suggest that the final products from the hydrogenation of the HO–CO complex have a purely statistical branching ratio. Therefore, HCOOH should be detected in the ice if CO₂ would be produced through this route, and this is not the case.

The oxidation of solid CO (dashed arrow) is also not likely to be the main formation reaction channel, since O₃, which would indirectly prove the presence of abundant O atoms in the ice, is not observed. O₃ ice has been detected in pure O₂ hydrogenation experiments only for temperatures higher than 25 K, when the penetration depth of the H atoms is higher than a few monolayers and the O₂ molecules are most likely more mobile (Ioppolo et al. 2010). The hydrogenation of O₃ ice (studied in Romanzin et al. 2011) is therefore not considered here. Furthermore, Roser et al. (2001) tentatively observed CO₂ formation through this channel only during the warm up of the ice and when CO molecules and O atoms were covered by a thick layer of H₂O ice, which allows the reactants to remain trapped in the ice for $T > 100\text{ K}$. The CO + O reaction contributes at best at high temperatures. The reaction HCO + O (dotted arrow) should also be ruled out, since HCO radicals prefer to react in a barrierless manner with H atoms forming H₂CO rather than with O atoms, as shown by Fuchs et al. (2009) and Ioppolo et al. (2011). Furthermore, as stated before, O atoms are not abundant in our ices.

At low temperatures, CO₂ is likely formed through the direct dissociation of the HO–CO complex in the ice (black arrow). The HO–CO complex is efficiently dissociated and, therefore, is not detected in our infrared spectra as a stable species. In a previous paper (Ioppolo et al. 2011), we observed this complex only in a water-rich environment. H-bonding should, indeed, improve coupling and heat dissipation through the ice, which stabilizes the HO–CO complex more easily in a polar than in an apolar environment. Our ice is mainly composed of CO and O₂, with a polar component on the surface of the ice. The amount of the HO–CO intermediate stabilized in the polar ice is also under the detection limit. Therefore, in a water-poor ice, the competition between dissociation and further

¹ Solid H₂CO₃ was observed only in control experiments by Ioppolo et al. (2011).

hydrogenation of the HO–CO complex is in favour of the dissociation.

H₂O is also formed through hydrogenation of the OH radicals (Fig. 2). Hence, the formation of CO₂ is linked to the formation of H₂O in the ice. This is consistent with the presence of CO₂ in polar (water-rich) interstellar ice mantles (see discussion in Section 5).

4.2 Hydrogenation of O₂ molecules

In this section, we investigate to what extent the hydrogenation of O₂, previously studied also for a pure ice, is affected by the presence of other molecules. Fig. 4 shows the H₂O₂ (top panels) and H₂O (bottom panels) column densities as a function of the H-atom fluence for the three different mixing ratios [CO:O₂ = 4:1 (circles), 1:1 (squares) and 1:4 (triangles)] and two temperatures investigated [15 K (left-hand panels) and 20 K (right-hand panels)]. For comparison, results from the hydrogenation of pure O₂ ice (Ioppolo et al. 2010) are also plotted (diamonds). Note that the top-right panel has a different scale for the column density axis than the other three diagrams. Formation rate and final yield of H₂O₂ and H₂O for all the investigated mixtures are lower than those from the pure O₂ ice hydrogenation. The differences in the final yield are more evident at higher temperature, where the yield for the mixed ices only moderately increases, whereas it increases with several monolayers for the pure O₂ experiments. This cannot be explained only by a low effective H-atom flux for the O₂ channel. Hence, the presence of CO in the mixture influences the final results. In a previous paper on CO + H we showed that H atoms can penetrate only a few layers of CO ice (Fuchs et al. 2009). Therefore, the presence of CO in the mixture most likely diminishes the penetration depth of H atoms into the ice compared to O₂ + H, and, therefore, desorption of H atoms from the ice can become important at higher temperatures. This explains the difference in the H₂O₂ and H₂O final yields compared to those from the pure O₂ ice, which increases with temperature.

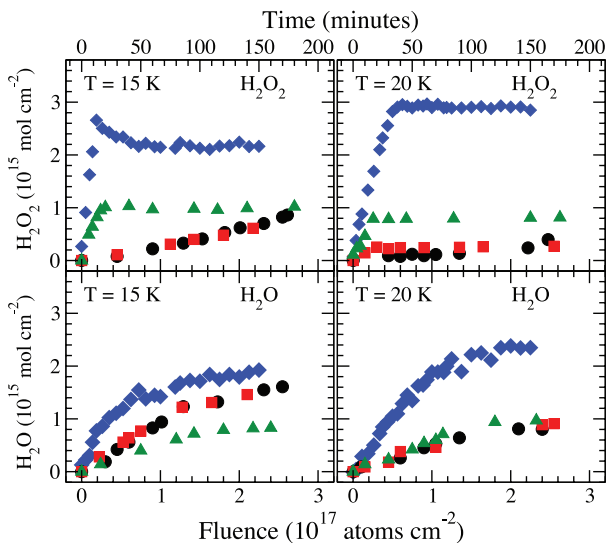


Figure 4. H₂O₂ (top panels) and H₂O (bottom panels) column densities as a function of the H-atom fluence and time of H-atom exposure at 15 K (left-hand panels) and 20 K (right-hand panels) for the three mixtures studied: CO:O₂ = 4:1 (circle), 1:1 (square) and 1:4 (triangle). For comparison, results from the hydrogenation of pure O₂ ice are plotted (diamond). Note the different vertical scale for upper right-hand panel.

The formation rates of H₂O and H₂O₂, which are reflected by the initial slopes of the curves, are also altered by the presence of CO in the ice. In the O₂-rich ice (1:4), the H₂O₂ column density shows the same behaviour as seen in pure O₂ ices: a constant formation rate is followed by a sharp transition towards saturation (Ioppolo et al. 2010). For high concentration of CO in the ice, the H₂O₂ column density increases with a much lower rate and does not appear to reach a steady state, even at the highest fluence. The H₂O₂ final yield increases with temperature, like for hydrogenation experiments of pure O₂ ice.

In the case of CO-rich ice (4:1), a more efficient conversion of H₂O₂ ice into H₂O ice can explain the high H₂O final yield with respect to the O₂-rich ice experiment. In a previous paper, we showed that H₂O₂ is more effectively formed in the bulk of the ice (Ioppolo et al. 2010). However, the presence of CO in the ice restricts the hydrogenation reactions to the surface of the ice. This means that a larger percentage of H₂O₂ formed at the surface of the ice is easily converted into H₂O. This may also explain the lower effective synthesis of H₂O₂ with the increase of the number of CO molecules in the ice. In addition, H₂O can be formed from OH radicals (see Fig. 1 and Cuppen et al. 2010), which can also react to form CO₂ in our ices. The H₂O column density is constant through almost all our experiments. This is also the case for CO₂ as shown in Section 4.1 and indicative for a correlation between the formation channels of these two species.

4.3 Hydrogenation of CO molecules

Fig. 5 shows the H₂CO (top panels) and CH₃OH (bottom panels) column densities as a function of the H-atom fluence for the three different mixing ratios [CO:O₂ = 4:1 (circles), 1:1 (squares) and 1:4 (triangles)] and two temperatures investigated [15 K (left-hand panels) and 20 K (right-hand panels)]. For comparison, results from the hydrogenation of pure CO ice (Fuchs et al. 2009) are also plotted (diamonds). For the hydrogenation of pure CO ice, the H-atom fluence is corrected according to recent H-atom flux measurements (Ioppolo et al. 2010), which improve the original H-atom flux

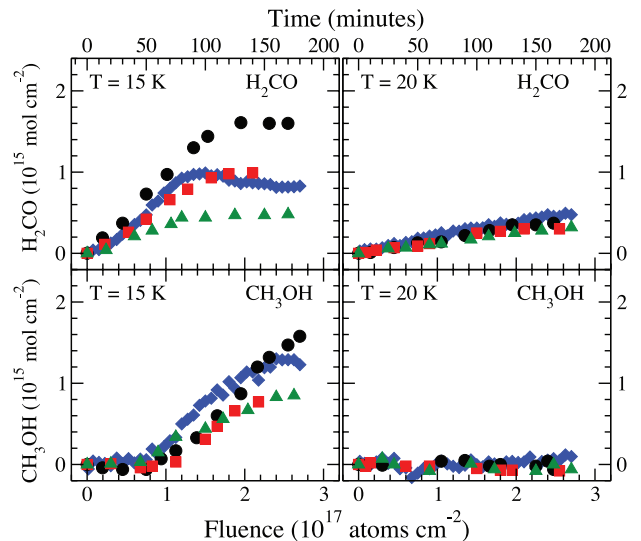


Figure 5. H₂CO (top panels) and CH₃OH (bottom panels) column densities as a function of the H-atom fluence and time of H-atom exposure at 15 K (left-hand panels) and 20 K (right-hand panels) for the three mixtures studied: CO:O₂ = 4:1 (circle), 1:1 (square) and 1:4 (triangle). For comparison, results from the hydrogenation of pure CO ice are plotted (diamond).

estimation derived in Fuchs et al. (2009). The hydrogenation of CO molecules in our mixtures shows the same behaviour seen for pure CO ice in terms of temperature dependence (Fuchs et al. 2009). An optimum in the final yield for H₂CO and CH₃OH is found at 15 K, while at 20 K no CH₃OH is formed and H₂CO has a low formation rate. The H₂CO and CH₃OH formation rates are hardly affected by the presence of O₂ in the ice (within the experimental uncertainties). The higher final yield for H₂CO and CH₃OH in the CO:O₂ = 4:1 experiment at 15 K compared to the pure CO ice cannot be explained by only a difference in the effective H-atom flux. In this case, the presence of O₂ in the ice, as a minor component, increases the penetration depth of H atoms in the ice compared to pure CO and, therefore, the probability that the H atoms get trapped and react in the ice. However, if the O₂ concentration is increased in the 15 K ice, the final yield decreases for both H₂CO and CH₃OH molecules. Clearly, the formation of H₂CO is more sensitive to the O₂ concentration in the ice than the formation of CH₃OH. As we saw for the hydrogenation of O₂ ice, the intermediate products (H₂O₂ and H₂CO, respectively) are more efficiently converted in the final products (H₂O and CH₃OH, respectively), when the ices are mixed.

4.4 Competition between the CO and O₂ channel

The results presented in the former sections reflect the competition between the two channels CO versus O₂, as shown in the left and right part of Fig. 1. It is clear from the experimental results that the presence of one component in the ice influences the reactivity of the other component. The formation rate of the CO hydrogenation reaction products is less affected by the presence of O₂ than the O₂ hydrogenation reaction products are affected by the presence of CO. This can be explained by the lower penetration depth of H atoms in CO ice and by the formation of CO₂ as an additional product, since OH radicals, formed through the O₂ channel, are used to form CO₂ instead of H₂O and H₂O₂.

In a CO-rich environment at 15 K, the presence of O₂ molecules enhances the production of H₂CO and CH₃OH, since H atoms can penetrate deeper in the ice than in the pure CO ice experiment. However, the formation rate of the final products seems not to be affected by the presence of the O₂ molecules in the ice. Moreover in a CO-rich ice, the formation of H₂O₂ is limited by the small amount of O₂ molecules in the ice, by the amount of OH radicals used to form CO₂ and by the lower penetration depth of H atoms in the ice, caused by the presence of CO molecules. H₂O₂ ice is also more efficiently converted to H₂O on the surface of the ice. This explains the high final yield for H₂O in a CO-rich environment at 15 K.

In an O₂-rich environment at 15 K, the formation of H₂CO and CH₃OH is limited by the small amount of CO molecules, which limits the formation of H₂O₂ and H₂O, since the penetration depth of H-atoms is lower than in the pure O₂ ice experiment.

At 20 K, the CO channel is not efficient, although the H atoms penetrate deep in the ice; at this temperature H atoms prefer to react with O₂ molecules. Also in this case, the final yields for H₂O₂ and H₂O are lower than those in the pure O₂ ice experiment.

5 ASTROPHYSICAL IMPLICATIONS

Observations by the *Infrared Space Observatory* (e.g. Gerakines et al. 1999; Nummelin et al. 2001; Gibb et al. 2004) and the *Spitzer Space Telescope* (e.g. Boogert et al. 2004; Bergin et al. 2005; Pontoppidan et al. 2005; Pontoppidan 2006; Whittet et al. 2007; Pontoppidan et al. 2008) have shown that roughly two-third of the

solid CO₂ observed in quiescent molecular clouds and star-forming regions is found in water-rich environments, suggesting that the formation routes of these two molecules are linked. The remaining CO₂ ice is predominantly found in a H₂O-poor, CO-rich environment (Pontoppidan et al. 2008). These observations suggest that the formation of CO₂ in dark quiescent clouds occurs in two distinct phases. In the early stages, CO₂ forms together with H₂O on the surface of the interstellar dust grains, creating a polar ice mantle. A second phase in the CO₂ formation occurs during the heavy freeze-out of CO. During this second phase, a H₂O-poor ice is formed.

Our experimental results indeed make it likely that CO₂ and H₂O are formed together in the early stages of the clouds through surface reactions assuming that both CO and OH are present in sufficiently high abundances. H₂O ice forms from continued hydrogenation of OH radicals formed on the surface of the dust grains. Alternatively, OH radicals can react with nearby CO molecules, which are present in small amounts in the ice before the strong CO freeze-out phase, forming CO₂ ice through the direct dissociation of the HO–CO intermediate. CO₂ can be also formed at low temperatures through the hydrogenation of the HO–CO complex, which can lead to the formation of HCOOH and H₂O + CO as well as CO₂ + H₂. The concentration of CO in the ice with respect to H atoms determines the probability of OH to react with a CO molecule or with another H atom.

In the second stage, during the heavy CO freeze-out, the gas density is $>10^5 \text{ cm}^{-3}$ and the CO accretion rate could be as high as, or even higher than, the H-atom accretion rate, which makes CO more abundant on the surface than H atoms. Any OH radicals will therefore more likely react with a nearby CO molecule than with H atoms. As long as some OH is present, CO₂ can thus be efficiently formed through the dissociation or further hydrogenation of the HO–CO complex, while just little H₂O ice is formed.

Energetic processing (UV irradiation and cosmic ray-induced photons) of polar and apolar ices is another efficient mechanism for CO₂ formation for specific (water-rich) environments (e.g. Hagen et al. 1979; Mennella et al. 2004, 2006; Loeffler et al. 2005; Ioppolo et al. 2009). All these channels, including the scheme discussed here, can contribute to the total CO₂ column density component observed in quiescent clouds.

In addition, in our experiments only a small amount of H₂O₂ ice is formed. This work shows that the competition between different channels (CO + H and O₂ + H), together with the penetration depth of the incoming H atoms into the ice, which decreases when the amount of CO in the ice increases, explains the differences in the H₂O₂ and H₂O final yield between a mixed and pure O₂ ice. Our results are consistent with the observed lack of H₂O₂ in the interstellar ices.

6 CONCLUSIONS

The experiments presented here show that CO₂ can be formed in the solid state for astronomically relevant temperatures via a thermal CO + OH reaction path. An efficient dissociation of the HO–CO complex is observed and this may explain the presence of CO₂ in polar and apolar interstellar ices in absence of UV irradiation. For the investigated laboratory conditions, CO₂ is formed efficiently and no strong dependency on temperature or ice composition is found. It is explicitly stated here that this conclusion does not rule out other CO₂ formation routes in space. Fig. 1, for example, shows how the H₂O formation through the O/O₂/O₃ + H channels is linked to the CO₂ formation. Here we investigated only the O₂ + H channel, and it should be noted that OH radicals can be also efficiently formed

on dust grains through the other H₂O formation channels as well as through the photodissociation of H₂O and CH₃OH ice.

The formation of CO₂ is linked to the formation of H₂O and, therefore, competes with the O₂ hydrogenation channel in our experiments. The competition of the two channels, together with the composition of the ice and the penetration depth of H atoms into the ice, explains the differences in the H₂O₂ and H₂O formation rate between a mixed and pure O₂ ice. Also the CO + H and O₂ + H channels are in competition and clearly affect each others final product yields. This is consistent with the lack of observed H₂O₂ in the interstellar ices. The formation rate for all the final products is found to be less sensitive on the mixture composition than the final yield. The penetration depth of the incoming H atoms is the main limiting factor. It depends on the composition of the ice and decreases when the amount of CO in the ice increases. Our results show that the formation rates found for H₂CO, CH₃OH, H₂O₂ and H₂O are similar within their experimental uncertainties to those found for the isolated CO and O₂ hydrogenation channels (corrected for the reduced effective H-atom fluxes). Those values, therefore, are still valid for use in astrochemical models. Finally, the experiments presented here chemically link CO₂ and H₂O, consistent with the observation of CO₂ in H₂O-rich environments in space.

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