Efficient Formation Pathway of Methyl Formate: The Role of OH Radicals on Ice Dust

A. Ishibashi®, H. Hidaka®, Y. Oba®, A. Kouchi®, and N. Watanabe®

Institute of Low Temperature Science, Hokkaido University N19W8, Kita-ku, Sapporo, Hokkaido 060-0819 Japan; watanabe@lowtem.hokudai.ac.jp

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Abstract

Three structural isomers of the C_2H_5O_2 molecule, namely, methyl formate (MF; HCOOCH_3), acetic acid (CH_3COOH), and glycolaldehyde (HOCH_2CHO), have attracted considerable attention as targets for understanding pathways toward molecular complexity in the interstellar medium (ISM). Among these isomers, MF is decisively abundant in various astronomical objects. For various formation pathways of MF, surface reactions on cosmic dust would play an important role. However, when compared to observations, the formation of MF has been found to be relatively inefficient in laboratory experiments in which methanol (CH_3OH)-dominant ices were processed by ultraviolet photons and cosmic-ray analogs. Here, we show experimental results on the effective formation of MF by the photolysis of CH_3OH on water ice at 10 K. We found that the key parameter leading to the efficient formation of MF is the supply of OH radicals by the photolysis of H_2O, which significantly differs from CH_3OH-rich experimental conditions. Moreover, using an ultrahigh-sensitivity surface analysis method, we succeeded in constraining the decisive formation pathway of MF via the photolysis of methoxymethanol (MM; CH_3OCH_2OH), which would improve our current understanding of chemical evolution in the ISM.

Unified Astronomy Thesaurus concepts: Dense interstellar clouds (371); Interstellar dust processes (838); Astrochemistry (75)

1. Introduction

Since the first detection by Brown et al. of methyl formate (MF; HCOOCH_3) in the star-forming region Sgr B2 (Brown et al. 1975), MF has also been detected in hot corinos (Bottinelli et al. 2007; Öberg et al. 2011) and outflow regions (Arce et al. 2008). Although a plausible formation process of MF in the gas phase is proposed recently (Taquet et al. 2016), the association reactions between radicals on the grain surface (Garrod & Herbst 2006) are considered as the important formation process of complex organic molecules (COMs) including MF. In recent years, MF was also found in cold molecular clouds (~10 K; Baczmann et al. 2012; Jiménez-Serra et al. 2016; Soma et al. 2018). Therefore, several formation mechanisms of MF have been theoretically proposed to explain the presence of abundant gaseous MF in cold molecular clouds: purely gas-phase formation (Balucani et al. 2015), gas-grain model using the chain-reaction mechanism on the grain surface (Chang & Herbst 2016), cosmic-ray-driven formation (Shingledecker et al. 2018), and nondiffusive formation on the grain surface (Jin & Garrod 2020). Those models were successful in explaining the abundance of MF fairly well. However, these are not still conclusive because the detailed information for building reliable models, such as reaction pathway, reaction rate constant, desorption rate by the chemical desorption, is insufficient.

Experimental studies of MF formation in the solid phase under the low-temperature conditions have been conducted mainly by the exposure of pure methanol solid to ultraviolet (UV) photons (Öberg et al. 2009; Paardekooper et al. 2016), because of a good correlation in astronomical observations between MF and CH_3OH (Yang et al. 2021). However, in laboratory experiments involving the photolysis of pure methanol solid, ethylene glycol (EG; HOCH_2CH_2OH) and ethanol (EtOH; CH_3CH_2OH) were produced in larger amounts than MF (Öberg et al. 2009; Paardekooper et al. 2016; Chuang et al. 2017), suggesting that the following reaction of photoproduced CH_2OH radicals dominates:

\[
\text{CH}_2\text{OH} + \text{CH}_3\text{OH} \rightarrow \text{HOCOCH}_2\text{CH}_2\text{OH(EG)},
\]

(1)

\[
\text{CH}_2\text{OH} + \text{CH}_3 \rightarrow \text{CH}_3\text{CH}_2\text{OH(EtOH)}. \]

(2)

On the one hand, the yields of MF and dimethyl ether (DME; CH_3OCH_3) were not significant, and these species would be derived from the following reactions:

\[
\text{CH}_3\text{O} + \text{HCO} \rightarrow \text{HCOOCH}_3\text{(MF),}
\]

(3)

\[
\text{CH}_3\text{O} + \text{CH}_3 \rightarrow \text{CH}_3\text{OCH}_3\text{(DME).}
\]

(4)

In addition, experiments involving a CO–methanol mixture show inefficient formation of MF and were unable to reproduce the observed predominance of MF over other COMs (Chuang et al. 2017).

These previous experiments were performed under CH_3OH-dominant conditions, which may be more relevant to the CO ice stage in dark molecular clouds (Cuppen et al. 2017). In ice mantles of grains, methanol would be present in H_2O-rich environments (CH_3OH/H_2O ~ 10%–30%; Boogert et al. 2015; Chu et al. 2020; Perotti et al. 2020). Recent experiments simulating the CO-ice stage demonstrated that a CO solid cannot fully cover H_2O ice surface. Instead, pure CO solid islands and isolated CO molecules form on water ice in the size range of dust grains (Kouchi et al. 2021a, 2021b). That is, the water ice is always exposed at the surface. On such grain surfaces, CO and its hydrogenated species can be adjacent to H_2O molecules each other. It is very likely that photochemical processes in such H_2O-rich sites of the surface differ from those in H_2O-poor site on pure CO solid, and thus there might be unknown processes leading to the effective formation of MF over other COMs. However, the experimental simulation of such processes suffers from technical difficulty in the detection of COM products and their precursors that may form under H_2O-rich, CH_3OH-poor conditions, where the yields of such
species would be most likely smaller than those obtained in pure methanol solid experiments. In the present study, we developed an ultrahigh-sensitivity surface analysis method, the so-called Cs\(^{+}\) ion pickup method, which enables us to mass analyze trace adsorbates, including radicals, on an ice surface with sensitivities 2 or 3 orders of magnitude higher than those possible with conventional methods, such as Fourier transform infrared spectroscopy (FTIR).

### 2. Experiments

#### 2.1. General Information about the Experimental Setup

As shown in Figure 1, the ion pickup experimental apparatus consisted of three parts; a main chamber, an ion source, and a UV light source. The typical pressure of the main chamber during UV irradiation was 10\(^{-5}\) Pa or less. An Al substrate located in the center of the main chamber was cooled to 10 K by a He refrigerator and heated to 320 K by an attached ceramic heater. The substrate was electrically insulated from the ground by a sapphire disk and thus could serve as a part of the lens system for ion transport upon application of an electric potential. Approximately 0.3 monolayers (ML; 1 ML = \(\sim10^{15}\) molecules cm\(^{-2}\)) of methanol, as an ice dust analog, was vacuum-deposited through a capillary plate on amorphous solid water (ASW) of 10 ML (1.5 ML in temperature-programmed pickup (TPP) experiments) pregenerated by background deposition of H\(_2\)O vapor on an Al substrate at 10 K. The amount of methanol was estimated by the same manner as our previous experiments (Hidaka et al. 2009). A quadrupole mass spectrometer (QMS) for the ion pickup measurement was installed at an angle of 45° to the substrate surface. At the head of the QMS instrument, the conventional ion source was removed, and instead, ring electrodes divided into two parts for assisting ion transport were installed. This specially designed ring ion lens played a key role in improving the detection sensitivity, which was \(\sim10^3\) times greater than the sensitivity of a conventional setup for the ion pickup method. The thickness and bulk composition of the sample solids were monitored by a reflection type FTIR.

The ion source, a Cs\(^{+}\) ion gun (Kimball Physics, ILG-6), was installed on the mirror surface side of the QMS at an angle of 45° with respect to the substrate surface. A Wien filter system was built in front of the Cs\(^{+}\) ion gun to remove other ion species contained in the Cs\(^{+}\) ion source. The ion source area was differentially pumped to prevent the flow of contamination to the main chamber.

A deuterium lamp (Hamamatsu Photonics: L7293), the wavelength of which was widely distributed across the UV region from 115 to 400 nm, was used as the UV light source. Because the spread of UV light from the lamp may cause emission of photoelectrons and degassing from the wall of the main chamber, the light path was surrounded by a cylindrical guide with a capillary plate on the top and differentially pumped inside. This differential pumping system at the UV light path effectively minimized undesired contamination on the sample surfaces. The average photon flux arriving at the substrate was \(\sim6 \times 10^{12}\) photons cm\(^{-2}\) s\(^{-1}\).

#### 2.2. Cs\(^{+}\) Ion Pickup Method

The basic idea of the Cs\(^{+}\) ion pickup method is well described elsewhere (Kang 2011). In the present experiments, low-energy (\(\sim17\) eV) Cs\(^{+}\) ions with a flux of \(\sim1\) nA picked up the adsorbed molecules (mass number M) on the outermost surface when they were scattered on the substrate. The adsorbates were detected by QMS as Cs\(^{+}\) ion–molecule complexes (mass number \(133 + M\)). The mass number M of the adsorbed species was identified by subtracting the mass number 133 of Cs from the detected mass number (Hang et al. 2001). This method allowed nondestructive detection of adsorbates on the surface, and the number of pickup adsorbates was sufficiently small to have no influence on the surface composition of the sample within the experimental duration. Therefore, continuous measurement was possible in real time (Appendix A1). Our developed method is much more
sensitive than conventional methods such as FTIR, as easily confirmed by comparing the detected molecules between the pickup spectrum and the IR absorption spectrum (see Appendix A2).

Because the electroconductivity of ice is low, the surface of ice can be charged by Cs\(^+\) ion injection. It was found that the degree of charging can change the detection efficiency of the ion pickup method. Therefore, the surface potential must be constant through any measurement. We controlled the voltage applied to the Al substrate to keep the surface potential constant. The surface potential, \(V_{\text{s}}\), is roughly described by the formula \(V_{\text{s}} = V_{\text{Al}} + V_{\text{charge}}\), where \(V_{\text{Al}}\) and \(V_{\text{charge}}\) represent the surface potential applied to the substrate and the potential achieved by charging, respectively. When the degree of charging changes on/in the ice sample, the surface potential can be maintained at a constant value by adjusting \(V_{\text{Al}}\). The criterion for the adjustment of \(V_{\text{Al}}\) is to keep the ion pickup signals at a maximum. This procedure is crucial for the experiments because the detection efficiency is extremely sensitive to the configuration of the electric potential between the sample surface and the ion detection systems. In fact, if the \(V_{\text{Al}}\) is not adjusted with the variation in the surface potential, the pickup signals rapidly disappear.

3. Results and Discussion

Using the Cs\(^+\) ion pickup method, we successfully detected a trace amount of photoproduct from methanol on ASW at 10 K. It is worth noting that photoproducts other than H\(_2\)CO were not detectable by FTIR but were detected by the Cs\(^+\) ion pickup method. The results will show the efficient formation of MF in consistent with astronomical observations.

Figure 2(a) shows the pickup mass spectrum of methanol adsorbed on ASW at 10 K before UV irradiation. A large peak at Mass 133 is due to the primary Cs\(^+\) ion. Peaks at Mass 151 (=133 + 18) and 165 (=133 + 32) originate from H\(_2\)O and CH\(_3\)OH, respectively (hereafter, only the mass of the picked-up species is denoted). For H\(_2\)O, molecules that are only loosely bonded to the surface are detectable, and thus the intensities do not reflect the surface number density of the total H\(_2\)O. The peak at Mass 33 was attributed to \(^{13}\)CH\(_3\)OH within the methanol reagent with the natural abundance ratio. In addition, peaks derived from simultaneous pickups of multiple H\(_2\)O and CH\(_3\)OH were detected (see Appendix A3). The spectrum of the sample after UV irradiation is shown in Figure 2(b). The peaks at Masses 30 and 28 correspond to photoproduced H\(_2\)CO and CO, respectively. In addition, OH, HCO (or \(^{13}\)CO) and CH\(_3\)OH (or H\(_3\)CO) radicals were detected at Masses 17, 29, and 31, respectively. Notably, new peaks appeared at Masses 60 and 62, corresponding to C\(_2\)H\(_4\)O\(_2\) and C\(_2\)H\(_6\)O\(_2\), respectively. The small peak observed at Mass 46 can be attributed to HCOOH rather than CH\(_3\)-derived DME and EtOH, as described in Appendix A4.

The peak at Mass 60 (C\(_2\)H\(_4\)O\(_2\)) was attributed to glycolaldehyde (GA) or MF, and that at Mass 62 was attributed to predominantly EG in previous works (Öberg et al. 2009; Paardekooper et al. 2016; Chuang et al. 2017), but other structural isomers were also possible for each mass: acetic acid (AA) at Mass 60 and methoxymethanol (MM; CH\(_3\)OCH\(_2\)OH) and dimethyl peroxide (DMP; CH\(_3\)OOCH\(_3\)) at Mass 62. To identify these molecules, the
pickup signals of the UV products at Masses 60 and 62 were measured while the temperature of the sample increased (hereafter referred to as the TPP spectrum), and the species can be identified by the temperatures at which the signals disappear, which correspond to the desorption temperatures of the target molecules (see Appendix A5). As seen in the TPP spectrum (Figure 3(a)), the signal at Mass 60 drops sharply at \( \sim 125 \) K. As a reference measurement, we obtained the TPP spectra of different samples in which gaseous MF was deposited on ASW (Figure 3(c)). The disappearance temperature of the peak at Mass 60 for the UV-photolyzed sample was consistent with that of MF for the reference measurement (see Figure 3(c)). As a result, the peak at Mass 60 is assigned to MF. Contributions from other isomers (GA and AA) are negligible since there was little signal above 150 K, which is below the desorption temperature of both molecules (\( \sim 160 \) K; Burke et al. 2015). For Mass 62, there are three main isomers: EG, DMP, and MM. As shown in Figure 3(b), the TPP signal at Mass 62 decreased sharply from \( \sim 180 \) K. The TPP spectra for reference measurements of MM and EG on the Al substrate were obtained (Figures 3(d) and (e)). The profile of the TPP spectrum for MM matches that for Mass 62 very well, while that for EG does not. Furthermore, the desorption temperature of DMP is known to be lower than that of MM (Zhu et al. 2019). Therefore, we conclude that the photoproduct corresponding to the peak at Mass 62 is mainly MM.

In the photolysis of pure methanol solid, the dominant product with a mass of 62 was EG, while MM was not positively identified (Öberg et al. 2009; Paardekooper et al. 2016), which implies that the formation of MM requires water molecules. We also deduce that the efficient formation of MF in the present experiment is correlated with MM formation. To gain more insight into the formation processes, variations in the radicals that may contribute to the formation of MF and MM were measured during UV irradiation of methanol on ASW at 10 K. As seen in Figures 4(a) and (b), with the consumption of methanol, CH\(_3\)O (or CH\(_2\)OH) radicals appeared first, followed by MM and, finally, MF.

Before discussing MF formation, the formation of MM, as a precursor of MF, is examined. As mentioned above, reactions that occur on the ASW surface should differ from those on the pure methanol solid. EG and MM are inferred to form by Reaction (1) and

\[
\text{CH}_3\text{O} + \text{CH}_2\text{OH} \rightarrow \text{CH}_2\text{OCH}_2\text{OH}(\text{MM}),
\]

respectively. In other words, the yields of MM and EG depend on the abundance ratio of the CH\(_3\)O and CH\(_2\)OH radicals on the surfaces. In the case of a pure methanol solid, CH\(_2\)OH-containing photoproducts such as ethylene glycol and ethanol were dominant comparing to the products originated from the CH\(_3\)O radical (Öberg et al. 2009; Paardekooper et al. 2016). From the latter results, the relationship CH\(_3\)OH > CH\(_2\)OH was deduced. In contrast, the formation of CH\(_2\)O leading to MM is expected to be enhanced on ASW. Here, the reactions involved in radical generation on ASW are shown below.
CH$_3$O formation (Xu & Lin 2007; Shannon et al. 2013; Antiñolo et al. 2016)

\[
H_2O + h\nu \rightarrow OH + H, \quad (6)
\]

\[
CH_3OH + OH \rightarrow CH_3O + H_2O. \quad (7)
\]

CH$_2$OH formation (Tachikawa 1993)

\[
CH_3OH + h\nu \rightarrow CH_2OH or CH$_3$O, \quad (8)
\]

\[
CH_3O + CH_2OH \rightarrow CH_3OH + CH$_2$OH. \quad (9)
\]

In the gas phase, Reaction (7) proceeds very fast due to the tunneling reaction at low temperatures (Xu & Lin 2007; Shannon et al. 2013; Antiñolo et al. 2016). On ASW, since H$_2$O is more abundant than methanol, it is expected that Reaction (7) occurs efficiently. That is, a new CH$_3$O formation pathway in addition to the photodissociation of methanol is possible on ASW, and thus the abundance ratio of CH$_3$O/CH$_2$OH tends to increase. In fact, as seen in Figure 4(a), the OH signal does not appear in the early stages of UV irradiation but appears when MM decreases (that is, MM production stops). This means that OH radicals are rapidly consumed by Reaction (7), supporting the above scenario (as a reference, see Appendix A6, for the behavior of OH on UV-irradiated pure ASW).

CH$_3$O generation can be enhanced by OH radicals, i.e., H$_2$O photodissociation. Because the photodissociation cross sections of H$_2$O and CH$_3$OH in the present UV region are comparable (Cruz-Díaz et al. 2014), the abundance ratio of CH$_3$O to CH$_2$OH can be reflected by the ratio of H$_2$O to CH$_3$OH on the outermost surface of the sample. In fact, the depletion rate of methanol for the photolysis of methanol on the ASW sample was approximately 2 times larger than that for pure methanol solid (Appendix A7), implying the occurrence of Reaction (7). In addition, the low detection of EG suggests that CH$_3$O > CH$_2$OH on ice surfaces (see Appendix A8).

Next, to confirm the pathway from MM to MF, MM deposited on the Al substrate was irradiated by UV photons. As shown in Figure 4(d), MF was found to be efficiently generated by the photoreaction of MM. Reactions leading to MF are thought to be

\[
CH_3OCH_2OH(MM) + h\nu \rightarrow HCOOCH$_3$(MF) + 2H(orH$_2$), \quad (10)
\]

together with Reaction (3). However, in the present experiment, because HCO is produced via H$_2$CO, its yield is thought to be small. In addition, the behaviors of HCO, H$_2$CO, and MF observed in Figures 4(a) and (c) do not seem to correlate with each other. Therefore, the contribution of HCO to MF formation would be minor under the present conditions.

Chang and Herbst reported that the enrichment of CH$_3$O density on grain surface by Reaction (7) enhances MF formation (Chang & Herbst 2016). Although it agrees with our result, our proposed formation mechanism of MF is different from that incorporated in their model. We propose the photodissociation of MM formed by the association of CH$_3$O + CH$_2$OH on grain surface as the additional formation pathway of MF. Besides photolysis and electron bombardments, Chuang et al. experimentally tested MF formation by simultaneous irradiation of CO and CH$_3$OH mixture with UV and H atoms at...
14 K (Chuang et al. 2017). They observed the enhancement of relative MF abundance to other major products such as EG and GA when comparing to the pure hydrogenation of the same sample. However, the relative abundance of MF obtained in their experiment was significantly lower than those of astronomical observations (see Figure 7 in Chuang et al. 2017). Consequently, the new reaction pathway proposed in the present study would be the key for the efficient process of MF formation on the cold solid surface.

4. Astrophysical Implications

The present results first show that MF is the overwhelmingly major photoproduct from methanol on ASW while neither EG nor GA are detected. As shown in Figure 5(a), the relative abundance of MF to its precursor, MM, in the present study has a good correlation with observations toward hot corinos of low-mass protostars (Jørgensen et al. 2018; Manigand et al. 2020) and high-mass star-forming regions (McGuire et al. 2017; El-Abd et al. 2019). This correlation would suggest that MF and MM are formed on dust grains in their parent clouds at low temperatures and released into the gas phase during warming up. In addition, the present results indicate that once trace amounts of methanol exist on ice mantles, COMs such as MF can be synthesized without the thermal diffusion of radicals in cold molecular clouds. Because HCO and CH3, both of which are minor products in the present experiment, would be abundant on realistic dust, Reactions (3) and (4) are also expected to be promoted, leading to the formation of MF and DME in cold molecular clouds (Figure 5(b)). This inference is consistent with the fact that the CH3O-derived products (MF, DME) were abundantly observed in various astronomical objects compared to the CH2OH-derived products (EG, GA; Herbst & van Dishoeck 2009; Oberg et al. 2009; Soma et al. 2018; Jørgensen et al. 2020). That is, the number density of CH3O on ice mantles should be high in cold molecular clouds, and such a situation is most likely brought by Reactions (6) and (7).

Occurrence of Reactions (6) and (7) requires coexistence of CH3OH with H2O on the mantle surface for a long period of time. This situation may not be consistent with the multilayered ice mantle model, where the H2O layer will be fully covered with CO and CH3OH mixed solid as time evolution progressed (Boogert et al. 2015). However, as the ice mantle, a more plausible structure has been recently found (Kouchi et al. 2021a, 2021b). Using the transmission electron microscope observation of ice mantle analogs produced in astrochemically relevant environments, the structure where CO-contained water ice and pure CO solids always coexist on the surface was confirmed.

In this study, we successfully reproduced the observed relative abundances of C2H4O2 isomers in cold molecular clouds for the first time. H2O was found to play an important role in COM formation from methanol. In other words, OH radicals are the key to forming various COMs, especially in cold molecular clouds. Our developed novel apparatus using the Cs+ ion pickup method can be further applied to other related studies, such as nitrogen-bearing COM formation, which is difficult by conventional analytical methods.
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Appendix

A1. Nondestructive Cs⁺ Ion Pickup Method

We performed experiments to ensure the nondestructive detection of CH₃OH. We monitored the masses of CH₃OH and possible ion-induced fragments (OH, CH₂O, and CH₂OH) from methanol on ASW without UV irradiation. No change was seen in CH₃OH, indicating that the pickup method had little effect on the surface density of CH₃OH. The intensities for CH₂O (or CH₂OH) and OH were under the detection limit. The intensity for H₂O increased because of the deposition of the residual H₂O gas in the chamber. In the pickup spectrum even after Cs⁺ ion injection for 3 hr (Figure 6(b)), CH₃ (Mass 148), MF (Mass 193), MM (Mass 195), etc., were not detected. Therefore, destruction of the sample by the Cs⁺ ion beam was almost negligible on the experimental timescale.

A2. High Sensitivity of the Cs⁺ Ion Pickup Method

To demonstrate the high sensitivity of the ion pickup method, we compared the FTIR measurements. Figure 7 shows the IR spectra of UV-photolyzed methanol on ASW under the conditions described in the main text. The amount of methanol that can be detected by the pickup method (Figure 2(b)) at approximately 1000 cps is approximately the detection limit (∼0.07 ML) of the FTIR measurements. H₂CO, which is produced in comparable amounts to the parent methanol, is expected to be detected at 1723 cm⁻¹, but it is hardly detected by FTIR due to its smaller absorption coefficient and overlap with the H₂O peak. MF is expected to appear near 1734 cm⁻¹ but cannot be detected because its yield is less than approximately 1/10 that of methanol. Accordingly, the pickup method is approximately 100 times more sensitive than FTIR, even for infrared-active species.

Figure 6. (a) Temporal signal changes in mass corresponding to the masses of CH₃OH, OH, H₂O, and CH₂O during Cs⁺ ion injection to methanol on ASW without UV irradiation. (b) Pickup spectrum after Cs⁺ ion injection for 3 hr.

Figure 7. (a) IR spectrum of UV-irradiated methanol (0.3 ML) on ASW (10 ML) for 3 hr. (b) Enlarged view of the 900–2000 cm⁻¹ region in (a).
A3. The Effect of Multiple-molecule Pickup in Mass Spectra

The mass peaks originated by a multiple-molecule pickup sometimes overlap with molecular species synthesized by the chemical reactions; for example, MF and two H₂CO pickup signals appear at the same mass number ($M = 60$). However, the origin of the mass peak can be easily distinguished in the same manner as molecular identification of the mass number containing isometric molecules described below (A5). In the case of $M = 60$, the intensities mainly decreased at the desorption temperature of MF but not at the temperature of formaldehyde. It indicates that the peak originates from MF. The mass peaks assigned as a multiple-molecule pickup in Figure 2 were also confirmed to originate from a multiple-molecule pickup by the TPP measurements.

A4. HCOOH (Mass 46) Formation from Methanol on ASW: No Products Derived from CH₃ Were Produced

There are three main structural isomers with a mass of 46. To identify the specific product corresponding to the peak at Mass 46, we obtained the TPP spectrum of this sample in UV-irradiated methanol on ASW (see Figure 8). The TPP intensities did not decrease at the desorption temperatures of DME and EtOH, indicating that both DME and EtOH are minor at the UV-irradiated sample surface. It is presumed that the reaction of the CH₃ radical did not occur on ASW, unlike on a pure methanol solid. The lack of CH₃ production on ASW also supports this hypothesis. This lack of reaction may be due to the low adsorption energy of CH₃ on ASW. Instead, the TPP intensities dropped significantly above the desorption temperature of HCOOH. This result clearly shows that the intensity of the peak at Mass 46 is governed by HCOOH. The enhancement of the intensity in the TPP profile above 130 K is thought to originate from the increase in pickup efficiency due to gradual removal of H₂O from the solid.

A5. Molecular Identification of the Mass Number Containing Isometric Molecules

Identification of molecular species with the same mass number was performed by measuring the TPP spectrum. In the TPP spectrum, the molecular species can be identified from the differences in the temperatures at which the pickup signals rapidly decrease because each adsorbed molecule has its own desorption temperature. We measured the reference TPP spectra of MF, MM, and EG to obtain each disappearance temperature. Measurements of the reference TPP spectra for MF and EtOH, which have lower desorption temperatures than H₂O, were conducted using samples that were prepared by molecular deposition on ASW at 10 K. For EG, which has a higher desorption temperature than H₂O, the reference TPP measurement was conducted without a H₂O layer sample formed at 165 K. MM was synthesized in the laboratory based on a previously reported protocol (Motiyenko et al. 2018). However, the synthesized MM sample contained large amounts of contamination (mainly methanol and H₂CO). Therefore, the MM sample for the reference TPP measurement was prepared by deposition on an Al substrate at 175 K, where H₂CO and CH₃OH hardly adsorb.

A6. Behavior of OH Radicals Formed by UV Irradiation of Pure ASW

When the methanol sample on ASW was continuously photolyzed, OH radicals were expected to form efficiently from the photodissociation of H₂O. However, the increase in OH was found to be slow, as shown in Figure 4(a). We ascribe this finding to the rapid OH consumption by Reaction (7), that is, $\text{CH}_3\text{OH} + \text{OH} \rightarrow \text{CH}_3\text{O} + \text{H}_2\text{O}$. If this hypothesis is true, the OH intensity should increase immediately when pure ASW is photolyzed. We measured the OH intensities for pure ASW during UV irradiation, as shown in Figure 9. The behavior of OH supports our hypothesis.

A7. Enhancement of the Methanol Attenuation Rate by Increasing the Number of ASW Layers

As discussed in the main text, if Reaction (7) enhances the destruction of CH₃OH in addition to direct photodissociation of CH₃OH, an increase in OH in the sample should enhance the attenuation rate of CH₃OH. Because OH radicals are formed by the photodissociation of H₂O, the number of H₂O molecules neighboring CH₃OH can be used to change the attenuation rate. We consider that the amount of deposited ASW can effectively control the yield of OH near CH₃OH. As seen in Figure 10, the attenuation rate of CH₃OH increases as the thickness (amount) of
ASW increases. This result clearly shows that H2O contributes to the enhancement of methanol destruction.

A8. Why DMP Is Not Detected despite the CH3O-rich Surface

On the CH3O-rich surface, DMP might be produced by the following reaction:

\[
\text{CH}_3\text{O} + \text{CH}_3\text{O} \rightarrow \text{CH}_3\text{OOCH}_3(\text{DMP}).
\]

However, O–O single bonds (peroxides) are generally known to be very fragile, perhaps resulting in photodecomposition into two CH3O radicals at a high rate (Toth & Johnston 1969). In addition, even if a large amount of CH3O is produced, it would be converted to CH2OH by Reaction (9).

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