Evidence of Surface Catalytic Effect on Cosmic Dust Grain Analogs: The Ammonia and Carbon Dioxide Surface Reaction

Alexey Potapov1, Patrice Theulé2, Cornelia Jäger1, and Thomas Henning3

1 Laboratory Astrophysics Group of the Max Planck Institute for Astronomy at the Friedrich Schiller University Jena, Institute of Solid State Physics, Helmholtzweg 3, D-07743 Jena, Germany; alexey.potapov@uni-jena.de
2 Aix-Marseille Université, CNRS, CNES, LAM, Marseille, France
3 Max Planck Institute for Astronomy, Königstuhl 17, D-69117 Heidelberg, Germany

Received 2019 April 16; revised 2019 May 28; accepted 2019 May 29; published 2019 June 11

Abstract

Surface chemistry on cosmic dust grains plays an important role in the formation of molecules at low temperatures in the interstellar and circumstellar environments. For the first time, we experimentally put in evidence the catalytic role of dust surfaces using the thermal reaction \( \text{CO}_2 + 2\text{NH}_3 \rightarrow \text{NH}_4^+\text{NH}_2\text{COO}^- \), which is also a proxy of radical–radical reactions. Nanometer-sized amorphous silicate and carbon grains produced in our laboratory were used as grain analogs. Surface catalysis on grains accelerates the kinetics of the reaction studied at a temperature of 80 K by a factor of up to 3 compared to the reaction occurring in the molecular solid. The evidence of the catalytic effect of grain surfaces opens a door for experiments and calculations on the formation of interstellar and circumstellar molecules on dust. Ammonium carbamate on the surface of grains or released intact into protostellar or protoplanetary disk phases can give start to a network of prebiotic reactions. Therefore, there should be a great interest to search for ammonium carbamate and its daughter molecule, carbamic acid, in interstellar clouds, protostellar envelopes, and protoplanetary disks.

Key words: astrochemistry – dust, extinction – methods: laboratory: solid state – molecular processes – techniques: spectroscopic

1. Introduction

Astrochemistry is a hot topic in the last decades due to the detection of complex organic molecules (COMs) in astrophysical environments, such as cold interstellar molecular clouds and hot protostellar molecular cores (see, for example, Kaitu et al. 1974; Bacmann et al. 2012; Vastel et al. 2014; Potapov et al. 2016), as well as in comets and meteorites (Cronin & Chang 1993; Elsila et al. 2007; Altwegg et al. 2017). One hypothesis about the source of organic compounds, which could serve as the basis of life on Earth, is their formation in the interstellar medium (ISM) and delivery to early Earth on board of meteorites (Oro 1961; Cronin & Chang 1993; Brack 1999; Pearce et al. 2017). Laboratory studies (Bernstein et al. 2002; Muñoz Caro et al. 2002; Nuevo et al. 2006, 2012; Elsila et al. 2007) have supported this idea and showed that spontaneous generation of amino acids and nucleobases in the ISM is possible in molecular ices covering the surface of cosmic dust grains.

Dust grains play a central role in the physics and chemistry of practically all astrophysical environments. They influence the thermal properties of the medium by absorption and emission of stellar light, provide surfaces for very efficient chemical reactions responsible for the synthesis of a major part of important astronomical molecules from simple ones such as \( \text{H}_2 \) and \( \text{H}_2\text{O} \) to COMs, and they are building blocks of planets. Functional groups and atoms at the surface of dust grains can participate directly in surface reactions and the grain surface itself has a threefold catalytic effect: (i) it provides a place where molecules can rapidly diffuse and react, (ii) it provides a third body to dissipate the energy released in exothermic bond formations, and (iii) it lowers the activation barriers of reactions. The first two effects have been shown by a vast number of laboratory experiments performed in interstellar ice analogs covering standard laboratory substrates such as gold, KBr, or HOPG, which are not characteristic of cosmic dust grains (for reviews see Theule et al. 2013; Linnartz et al. 2015; Öberg 2016). There is only a handful of studies demonstrating the formation of \( \text{CO}, \text{CO}_2, \) and \( \text{H}_2\text{CO} \) molecules involving surface atoms of carbonaceous grains (see Potapov et al. 2017 and references therein). The same applies to the catalytic effect of dust surfaces on surface reactions in ices or on bare grains. Only a few studies were performed on the formation of molecular hydrogen and water on silicate surfaces (see Gavilan et al. 2014; He & Vidali 2014 and references therein). Water ice can also participate in the sequence of elementary reactions leading to the formation of COMs by lowering one or more reaction barriers (Danger et al. 2014; Noble et al. 2014).

To evaluate the catalytic effect of the dust surface we use the model reaction \( \text{CO}_2 + 2\text{NH}_3 \rightarrow \text{NH}_4^+\text{NH}_2\text{COO}^- \) in the submonolayer regime. The two reactants, \( \text{CO}_2 \) and \( \text{NH}_3 \), are two of the several main components of interstellar and cometary ices (Tielens 2013). The reaction is purely thermal and leads to the formation of ammonium carbamate, \( \text{NH}_4^+\text{NH}_2\text{COO}^- \), and carbamic acid, \( \text{NH}_2\text{COOH} \), at temperatures higher than 70 K (Bossa et al. 2008; Noble et al. 2014; Ghesquiere et al. 2018).

2. Experimental Part

The experiments have been performed using the new ultra-high vacuum (UHV) setup, INterStellar Ice Dust Experiment (INSIDE), recently developed in Jena and presented in detail elsewhere (Potapov et al. 2019). The base pressure in the main chamber of the setup is a few \( 10^{-11} \text{ mbar} \) allowing for the reproduction of the physical conditions in dense interstellar clouds and the performance of clean experiments without
additional adsorption of species such as water from the chamber volume.

In brief, nanometer-sized, amorphous carbon ($^{13}$C) and silicate (MgFeSiO$_4$) grains were produced in a laser ablation setup (Jäger et al. 2008; Potapov et al. 2018) by pulsed laser ablation of a graphite or Mg:Fe:Si target and subsequent condensation of evaporated species in a quenching atmosphere of a few mbar He and H$_2$ for carbon grains or He and O$_2$ for silicate grains. Condensed grains were deposited onto a KBr substrate. The thickness of the deposit was 150 nm for MgFeSiO$_4$ (density 3.7 g cm$^{-3}$) and 70 nm for carbon (density 1.7 g cm$^{-3}$) controlled by a microbalance (sensitivity 0.1 nm) using known values for the deposit area and density. A reference surface of pure KBr was also used.

After the deposition of grains, the samples were extracted from the deposition chamber (exposed to air), fixed on a sample holder, and inserted into the pre-chamber of INSIDE, where they were annealed at 200°C during two hours to remove possible surface contamination by molecular adsorbates. After annealing, the sample holder was moved to the UHV chamber of INSIDE and fixed on the cold head. The pure KBr substrate used in this study was also annealed. NH$_3$ and CO$_2$ (purity 5.5) with a ratio of 4:1 were deposited on the surfaces of KBr, $^{13}$C grains, or MgFeSiO$_4$ grains at 15 K. The gases were introduced through two separated gas lines. The CO$_2$ and NH$_3$ ice thicknesses were calculated from their vibrational bands at 2342 and 1073 cm$^{-1}$ using the band strengths of 7.6 x 10$^{-17}$ cm molecule$^{-1}$ (Gerakinus et al. 1995) and 1.7 x 10$^{-17}$ cm molecule$^{-1}$ (d’Hendecourt & Allamandola 1986), and assuming a monolayer column density of 1 x 10$^{13}$ molecules cm$^{-2}$ and a monolayer thickness of 0.3 nm.

After the deposition of ice, the samples were heated up to 80 K with a heating rate of 100 K minute$^{-1}$, the initial time was set to zero, and the isothermal kinetics experiments were performed. The temperature was chosen as high enough to perform the efficient reaction and low enough to prevent an active desorption of CO$_2$ and NH$_3$. The formation of the NH$_4^+$-NH$_2$COO$^-$ product was monitored by the growth of its characteristic vibrational bands at 1550 and 1385 cm$^{-1}$ as a function of time. The column density of NH$_4^+$-NH$_2$COO$^-$ was calculated from the 1550 cm$^{-1}$ band using the band strength of 2 x 10$^{-17}$ cm molecule$^{-1}$ (Bossa et al. 2008). After the kinetics experiments were completed, temperature-programmed desorption (TPD) experiments with a heating rate of 2 K minute$^{-1}$ were realized in the temperature range of 80–300 K. Infrared (IR) spectra were measured using an FTIR spectrometer (Vertex 80v, Bruker) in the transmission mode. Mass spectra were measured by a quadrupole mass spectrometer (HXT300M, Hositrad) equipped with a multiplier detector (multiplication factor of 10$^3$). We followed CO$_2$ on m/z 44, NH$_3$ on m/z 17, and ammonium carbamate on m/z 61 and 78.

The morphology of carbon and silicate grains can be understood as a porous layer of rather fractal agglomerates. The very high porosity of grains (Sabri et al. 2014) means an existence of a very large surface. In Figure 1, we show a high-resolution electron microscopy image of amorphous silicate grains, where a high porosity and a large surface are clearly observable. The inset shows carbon grains deposited on a KBr substrate with a nominal surface area of 1 cm$^2$. It was recently shown that 130 nm of water ice mixed with carbon grains and deposited on a nominal substrate surface of 1 cm$^2$ corresponding to a theoretical coverage of about 400 monolayers (ML) desorb following the first-order kinetics of desorption (Potapov et al. 2018). This result was explained by the desorption of water molecules from a large surface of clusters composed of carbon grains. Therefore, the real surface of dust grains deposited on a substrate can be two orders of magnitude larger than the nominal substrate surface of 1 cm$^2$.

To investigate the effect of the surface, it is important to know the monolayer–multilayer transition on the porous carbon and silicate grains we use. The thickness corresponding to the monolayer–multilayer transition is given by the transition between the zeroth-order and first-order form of the Polanyi–Wigner equation (Polanyi & Wigner 1928). According to our TPD curves for ammonia and carbon dioxide taken after the kinetics experiments were completed, we can conclude that the monolayer–multilayer transition takes place on the grains at the ice thickness (calculated from the vibrational bands of CO$_2$ and NH$_3$ as described above) about 200 nm.

3. Results

In Figure 2, one can see an example of the IR spectra taken after the deposition of a CO$_2$:NH$_3$ mixture at 15 K and after 4 hr of the isothermal kinetics experiment at 80 K. A detailed assignment of the IR absorption bands of NH$_3$:CO$_2$ ices is done elsewhere (Bossa et al. 2008; Noble et al. 2014). Two bands related to NH$_4^+$NH$_2$COO$^-$ at 1550 and 1385 cm$^{-1}$ are immediately observable in the IR spectra after heating of the samples up to 80 K.

An example of the results of isothermal kinetics experiments at 80 K is shown in the inset of Figure 2. The exponential decay of CO$_2$ and the corresponding growth of NH$_4^+$NH$_2$COO$^-$ exhibit pseudo-first-order reaction kinetics (Noble et al. 2014). For the determination of the rate coefficient we followed the procedure described in Noble et al. (2014), solving the kinetic equations

\[
(CO_2)(t) = (CO_2)_0 \times e^{-kt}
\]

\[
(NH_4^+NH_2COO^-)(t) = (CO_2)_0 \times (1 - e^{-kt}).
\]

Reaction-rate coefficients obtained for different surfaces and ice thicknesses are presented in Table 1 and in Figure 3. The reaction rate coefficients presented were obtained from
isothermal curves following the formation of ammonium carbamate. Using isothermal curves following the CO₂ signal, the reaction rates on grains for ices thinner than 200 nm were found to be higher (∼10%) than the rates obtained from the ammonium carbamate curves indicating a slight desorption of CO₂ (mainly trapped in the NH₃ matrix) in thin ices on grains.

After the completion of the kinetic studies, TPD measurements in the temperature range of 80–300 K provided signals at masses (m/z) 44 and 17 corresponding to the desorption of CO₂ and NH₃ at about 95 and 105 K and signals at masses 61 (NH₃COOH), 44 (CO₂), and 17 (NH₃) related to the desorption and fragmentation of NH₄⁺NH₂COO. The signal at mass 78 corresponding to the intact NH₄⁺NH₂COO⁻ molecule was not observed. In Figure 4, we present TPD curves for NH₄⁺NH₂COO⁻ obtained from the mass spectra by taking the signal at mass 61 and from the IR spectra by taking the first temperature derivative of the integrated intensity of the 1550 cm⁻¹ band. The curves were measured with a ramp rate of 2 K minute⁻¹ for 200 nm ice on carbon grains after 4 hr of the isothermal kinetic experiment. The results are in agreement with previous experiments (Noble et al. 2014).

4. Discussion

The reaction rate coefficients obtained in the present study are equal for all surfaces when the ice thickness is about 500 nm. The reaction rate coefficient for the reaction on KBr stays constant in the limits of uncertainties in the whole range of ice thicknesses. However, with the decrease of the ice thickness, starting from 200 nm thickness, we observed a dramatic increase of the reaction rate coefficient reaching values close to 1 × 10⁻³ s⁻¹ at a nominal ice coverage of about 35 nm or 100 ML. Thus, we measured up to three times higher reaction rates on grains compared to the molecular solid on KBr depending on the ice thickness. If we compare with the literature value 4.9 × 10⁻⁵ s⁻¹ (Noble et al. 2014) measured at 80 K for a NH₃:CO₂ 4:1 mixture in a molecular solid, the reaction rate on grains is up to 20 times higher. The difference between our KBr values and the literature value could be explained by different experimental procedures of two studies. The increased reaction rates for grains compared to those measured for NH₃:CO₂ molecular solids can be explained in
two ways: (i) an increased diffusion rate of species and (ii) a participation of dust surfaces in the reaction studied. The first explanation is purely physical: at low temperature the reaction-diffusion is diffusion-limited. The diffusion of the reactant molecules on a surface is much faster than in a bulk material (Mispelaer et al. 2013; He et al. 2018). It has been shown that the diffusion of CO2 and NH3 reactants in the volume of ice occurs along the surface of cracks generated by the structural changes in water ice mantles upon crystallization, which leads to a dramatic increase of the production rate of ammonium carbamate as compared to what it can be with bulk diffusion (Ghesquiere et al. 2018). In our case of a pure NH3:CO2 solid, the surface is provided by aggregates/clusters of nanometer-sized carbon and silicate grains seen in Figure 1.

The second explanation implies a participation of the dust surface in the CO2 + 2NH3 reaction. First, the reaction CO2 + 2NH3 + dust can lead to the formation of an intermediate weakly bound CO2−NH4 complex, which could be stabilized by a third body (dust surface) that helps to overcome the reaction barrier. The formation of weakly bound complexes on the surface of grains in the ISM is a matter of a number of experimental and theoretical studies (see Potapov 2017 for a brief review). Second, the reaction barrier of the CO2 + 2NH3 reaction can be different for different surface compositions. It can vary by several kJ mol−1 depending on the concentration ratio of the NH3:CO2:H2O cluster (Noble et al. 2014). To the best of our knowledge, no calculations on carbon or silicate surfaces exist.

5. Astrophysical Implication

Submonolayer reactions are important in two astrophysical stages or environments: (i) during the transition from diffuse to dense clouds and the formation of the first monolayer, which usually takes place during the first 103 yr; and (ii) during the desorption of the volatile mantles and the subsequent concentration of refractory materials taking place at the snowline of protostellar envelopes and protoplanetary disks. The fact that the surface has an effect on the submonolayer or on the bottom most layer can result in an enhanced solid-state reactivity for reactions involving molecules and/or radicals.

The reaction products of NH3 and CO2, ammonium carbamate and carbamic acid, can store CO2 and NH3 on the surface of dust grains in interstellar clouds. During the protostellar phase, grains are heated and above 250 K; when the carbamic acid and ammonium carbamate desorb, CO2 and NH3 can be introduced into the gas phase at much higher temperatures than their normal desorption temperatures.

Once delivered on our planet, both ammonium carbamate and carbamic acid can further evolve under the primitive Earth environment. It was discovered that ammonium carbamate can be converted into urea CO(NH3)2 and water in a temperature range between 135°C and 200°C (Clark et al. 1933). Formamidine and urea were also produced in interstellar ice analogs by UV irradiation of a HNCO ice (Raunier et al. 2004) or a CH3OH:NH3 ice mixture (Nuevo et al. 2010). We can assume that UV irradiation of solid ammonium carbamate may also lead to the formation of urea. Urea is a molecular corner stone in the search for the origin of life (Miller & Urey 1959). It can be a precursor of pyrimidine required for the synthesis of nucleobases in RNA molecules (Robertson & Miller 1995). Urea was tentatively detected in the ISM, on grains as well as in the gas phase (Raunier et al. 2004; Remijan et al. 2014).

Thus, ammonium carbamate on the surface of grains or released intact into the protostellar or protoplanetary disk phases can give start to a network of prebiotic reactions. The search for ammonium carbamate and its daughter molecule, carbamic acid, in interstellar clouds, protostellar envelopes, and protoplanetary disks should be of great interest. However, the problem is that these molecules are not stable at room temperature in the gas phase. Therefore, high-resolution gas-phase spectra of the molecules, which allow a radio-astronomical search, are not available. One possibility to obtain their spectroscopic passports is to synthesize the molecules in interstellar ice analogs and detect them above the surface of ice immediately after their release into the gas phase. This can be realized by a combination of solid-state chemistry and high-resolution gas-phase spectroscopy in one experiment. Such an idea is currently under development in a few groups worldwide.

There are a number of important surface thermal reactions for prebiotic astrochemistry, for example NH4+ + HCN → NH4+CN− (Noble et al. 2013) and CO2 + 2CH3NH2 → CH3NH2CHNCO− (Bossa et al. 2009). NH4+CN− can react with CH3NH2, which is a product of HCN hydrogenation, leading to aminoacetonitrile NH2CH2CN that can form glycine according to the Strecker synthesis. CH3NH2+CHNCOO− acts as a glycine salt precursor in UV environments. In this context, it is also worth mentioning hydrogenation reactions such as the hydrogenation of CO leading to the formation of methanol (Hiraoka et al. 1994) and the hydrogenation of HCN leading to the formation of methylamine (Theule et al. 2011), a direct precursor of glycine. All of these reactions can occur on a much shorter timescale than it is thought now.

6. Summary

The present study demonstrates that the rate coefficients for the thermal reaction CO2 + 2NH3 → NH4+NH2COO− on the surface of nanometer-sized carbon and silicate grains, which are reliable analogs of interstellar and circumstellar dust grains, are up to three times higher compared to the rate coefficients measured in the molecular solid. Thus, the catalytic effect of grains is clearly observed and must be taken into account. This result shows the role of the surface in the kinetics of solid-state reactions. A faster kinetics, on this particular reaction or other ones, especially radical–radical reactions, should have a great impact on astrochemical modeling involving surface chemistry.

We would like to thank two anonymous referees for questions, suggestions, and corrections that helped to improve the manuscript. This work was partly supported by the Research Unit FOR 2285 “Debris Disks in Planetary Systems” of the Deutsche Forschungsgemeinschaft (grant JA 2107/3-1).

ORCID iDs
Alexey Potapov @ https://orcid.org/0000-0002-3699-7477

References
Altwegg, K., Balsiger, H., Berthelier, J. J., et al. 2017, MNRAS, 469, S130
Bacmann, A., Taquet, V., Faure, A., Kahane, C., & Ceccarelli, C. 2012, A&A, 541, L12
Bernstein, M. P., Dworkin, J. P., Sandford, S. A., Cooper, G. W., & Allamandola, L. J. 2002, Nature, 416, 401
Bossa, J. B., Duvernay, F., Theule, P., et al. 2009, A&A, 506, 601
Noble et al. 2013
Robertson et al. 2017
