

Hydrogen addition reactions of simple aliphatic hydrocarbons (C₂H₂ and C₂H₄) ~ clue to the formation mechanism of cometary C₂H₆~

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Comets are thought as remnants of planetesimals formed in the early solar nebula. Their nuclei consist of volatile ices and dust grains that were formed in the pre-solar molecular cloud and probably chemically altered in the proto-planetary disk. Although the outline of evolution from a molecular cloud to the disk is basically understood, detailed chemical processes are still in debate. To investigate the links between proto-planetary disk and the molecular cloud, we focused on cometary ethane (C₂H₆) which is considered to form through the hydrogen addition reactions on the cold grain surfaces. C₂H₆ and acetylene (C₂H₂), the parent molecule of the C₂H₆, have been observed in multiple comets since 1996 and their abundances relative to water (the major component of cometary ices) is ~0.001 with some variations [1]. Such variation might be caused by the difference in the mixing ratios between the materials originated in the molecular cloud and the disk-processed materials. C₂H₆ has never been detected in molecular cloud thus we evaluate experimentally the hydrogenation processes to form the C₂H₆ from C₂H₂ (C₂H₂ → C₂H₃ → C₂H₄ → C₂H₅ → C₂H₆). In the previous experimental studies [2], it was concluded that the reaction from C₂H₄ to C₂H₆ is considerably rapid than the reactions from C₂H₂ to C₂H₄ and it would be a reason for the absence of C₂H₄ in comets. To investigate those reactions quantitatively in realistic conditions of molecular clouds, we performed the laboratory measurements of hydrogen addition reactions of C₂H₂ and C₂H₄ on amorphous solid water.

The experiments were conducted by using laboratory setup for surface reaction in interstellar environment (LASSIE) at Institute of Low Temperature Science, Hokkaido University [3]. Atomic hydrogen used for the reactions were produced by the dissociation of hydrogen molecules in micro-wave induced plasma. The kinetic temperature of hydrogen atoms were ~120K and the H-atom flux was ~10¹³ cm⁻²s⁻¹. The samples of pure solid C₂H₂, C₂H₄ and those on ASW were produced on the substrate located in the center of the main chamber at 10K and 20K. Infrared absorption spectra of the ices were measured by FTIR before and during the exposure of H-atom.

Although our measurements show basically the same trend as shown in the previous studies, the difference in reaction rates for C₂H₂ and C₂H₄ was found to be much less than that previously reported. We will discuss the temperature and thickness dependence of the time constant for the sample ices in the poster.

References

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