

SUPPRESSED RADIOLYSIS OF HYDROGEN PEROXIDE IN WATER ICE - HYDROGEN MIXTURES.

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Introduction: Water ice has been identified as a major constituent of icy surfaces in the outer solar systems / interstellar medium, where it is bombarded by solar radiation, solar wind ions, cosmic rays, energetic particles trapped in the planetary magnetospheres and micrometeorites.

Radiation chemistry leads to the formation of species such as H, O and OH within the ice films from dissociation of water molecules due to ion bombardment. Oxygen enrichment occurs when the H atoms are preferentially lost to vacuum, due to its small size which favors the diffusion and desorption of H out of the ice films. The OH radicals, on the other hand, are much less mobile compared to the H atoms due to the cage effect, which tend to confine the OH radicals within the vicinity of the position from where they are formed, or the ion track. Most of the OH will recombine with H to form H₂O and the excess OH radicals can combine with OH radicals formed in the vicinity by a following ion or those diffused from nearby ion tracks, to form Hydrogen Peroxide (H₂O₂). The H₂O₂ production is found to decrease with increasing temperature, and is attributed to the competing phenomena of OH diffusion away from the ion track and hydrogen escape from the ice [1], so that H₂O₂ production at higher temperatures is less efficient due to higher H escaping rate.

Experiments of photolysis of H₂O₂ in ice films with 193 nm photons at various ambient O₂ pressures[2] showed that the oxygen enrichment in the irradiation environment enhances the H₂O₂ production. Radiolysis of ice films with 50 keV protons at low ambient O₂ pressure showed no significant enhancement of H₂O₂ production for fluences that saturates O₂ trapping due to ice compaction [3], a fluence that is lower than required to saturate H₂O₂ production. However, higher ambient O₂ pressures and higher total fluences might result in enhancement of H₂O₂ production, probably by transferring trapped O₂ molecules into H₂O₂ through radiation chemical reactions.

Inspired by the above oxygen enrichment experiments, we tried a different method of enriching hydrogen in the ice films by growing an ice + H₂ mixture, and studying H₂O₂ production under bombardment of 100 keV H⁺ with H₂ ambient pressure in the vacuum chamber.

The Experimental Setup: The experiments were conducted in a UHV chamber with a base pressure of 10⁻¹⁰ mbar. Water ice films were vapor deposited onto a Li-He cooled gold coated quartz crystal microbalance

at 20 K, through a collimating micro-capillary array doser. The areal mass of the deposited films was measured by converting the change of the resonant frequency of the microbalance due to mass increase to column density. The ice films were irradiated at 9 degree incidence with a mass-analyzed collimated 100 keV H⁺ beam from a 300 keV ion accelerator. The reflectance spectra of the ice films within the region of 650 - 9000 cm⁻¹ were collected at an incident angle of 35 degree with a Thermo-Nicolet Nexus 670 Fourier Transform infrared spectrometer (FTIR) at 2 cm⁻¹ resolution. The reflectance R was obtained by dividing the reflectance spectra by the spectrum of the gold mirror, and then converted to optical depth, -ln(R). Methods for quantification of the infrared spectra are published elsewhere [4].

Results: In the first series of experiments, we grew pure ice films at 20 K at 45 degree incidence angle to enhance the porosity of the film, in order to enhance the adsorption of H₂ with an ambient pressure. We maintained constant ambient H₂ pressures between 3 × 10⁻⁸ and 3 × 10⁻⁵ Torr. Then we introduced 100 keV H⁺ ions into the ice film (flux: $f = 1 \times 10^{12} \text{ cm}^{-2}\text{s}^{-1}$). The production of H₂O₂ was monitored by the intensity of the H₂O₂ absorption band at ~ 2860 cm⁻¹. We found that the H₂O₂ band area is not affected consistent with blank experiments without ambient H₂.

In comparison, we repeated the experiments with H₂O + H₂ mixtures instead of pure ice films. During the growth of the mixture, the ambient H₂ pressure was first introduced and maintained at a constant value, then H₂O vapor was leaked into the chamber. During the proton irradiation (100 keV H⁺, flux: $f = 1 \times 10^{12} \text{ cm}^{-2}\text{s}^{-1}$), the ambient H₂ pressure was kept constant. We varied the ambient pressure during the growth of the ice mixtures and repeated the above procedures. Surprisingly, we found that at an ambient pressure $P > 3 \times 10^{-6}$ torr, the amount of H₂O₂ produced is negligible and the hydrogen enrichment leads to suppression of H₂O₂ production. For pressures $P < 3 \times 10^{-7}$ torr, we observed a partial suppression of H₂O₂ production. We explored the flux dependence of H₂O₂ production under ambient H₂ pressure (3 × 10⁻⁷ torr) and did not find a further suppression of H₂O₂ production for the lowest ion fluxes we used ($f = 5 \times 10^{10} \text{ cm}^{-2}\text{s}^{-1}$).

We investigated the temperature dependence. At 22 K, we repeated the experiments at ambient H₂ pressures upto 3 × 10⁻⁶ torr and did not observe a suppression of H₂O₂ production, suggesting that lower temperature favors the hydrogen enrichment that lead to

suppression of H_2O_2 production, consistent with the reported findings [1] of lower H_2 escape rate from ice films at lower temperatures leading to higher H_2O_2 production.

Discussion: The results of this research showed that the H_2O_2 production in ice + H_2 mixture is suppressed compared to pure ice films, and the suppression is stronger at lower temperatures and higher ambient H_2 pressures. In the interstellar medium, ~99% of the mass are attributed to gas phase matter, ~3 quarters of which are H/H_2 , ice rich grains may trap H/H_2 in the lattice when they are formed, Environmental radiation of these hydrogen enriched grains may not be able to produce as much H_2O_2 as in pure ice grains.

References:

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