

Abstract Submitted  
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**H<sub>2</sub>-Assisted Ternary Recombination of H<sub>3</sub><sup>+</sup> with Electrons at 300 K**<sup>1</sup> RAINER JOHNSEN, University of Pittsburgh, PETR DOHNAL, PETER RUBOVIC, ABEL KALOSI, MICHAL HEJDUK, RADEK PLASIL, JURAJ GLOSIK, Charles University Prague — Afterglow measurements in ionized He/Ar/H<sub>2</sub> gas mixtures at 300 K show that the recombination of H<sub>3</sub><sup>+</sup> ion with electrons is very strongly enhanced in the presence of molecular hydrogen. In the experiments the decay of H<sub>3</sub><sup>+</sup> ions was measured by near-infrared (NIR) absorption spectroscopy (SA-CRDS).<sup>2</sup> Rather surprisingly, the H<sub>2</sub>-assisted three-body recombination coefficient ( $K_{\text{H}_2} = (8.7 \pm 1.5) \times 10^{-23} \text{ cm}^6\text{s}^{-1}$ ) exceeds by more than two orders of magnitude the corresponding He-assisted coefficient ( $K_{\text{He}} = (3.3 \pm 0.7) \times 10^{-25} \text{ cm}^6\text{s}^{-1}$ ) that we measured earlier.<sup>3</sup> Formation of faster recombining H<sub>5</sub><sup>+</sup> cluster ions does not play a significant role at temperature near 300 K. The ternary processes are found to saturate at high He and H<sub>2</sub> densities, suggesting that recombination proceeds by a two-step process, electron capture (with a rate coefficient  $\alpha_F = (1.5 \pm 0.1) \times 10^{-7} \text{ cm}^3\text{s}^{-1}$ ) into a long-lived Rydberg state with an excited core, followed by collisional stabilization. While these findings provide a plausible explanation for some of the discrepancies between earlier afterglow measurements of H<sub>3</sub><sup>+</sup> recombination, the exact nature of these long-lived complexes, and their collisional interactions remain to be elucidated.

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<sup>2</sup>P. Macko et al, *Int. J. Mass Spectrom.* **233**, 299 (2004).

<sup>3</sup>R. Johnsen et al, *J. Phys. Chem. A* **117**, 9477 (2013).

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