HIGH-RESOLUTION INFRARED SPECTRA OF THE OH-STRETCHING BANDS OF PROTONATED WATER DIMER, $H_5O_2^+$

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We present high-resolution infrared action spectra of cold $H_5O_2^+$. For this purpose the mass selected parent ions are stored in a cryogenically cooled 22-pole ion-trap (COLtrap). There we employ a two-color-photodissociation scheme where first the symmetric or the anti-symmetric O-H-stretching band is excited by a narrow linewidth cw-OPO. Then, Light from a CO₂ laser is used to efficiently dissociate the parent molecule. The infrared-absorption of the parent ion is recorded by the appearance of H_3O^+ photoproducts. This procedure follows the seminal approach first invented in the group of Y.T. Lee ^{*a,b*}. The rotationally resolved and basically background-free spectrum exhibits a complex structure, making the assignment of individual ro-vibrational tunneling features challenging. Nonetheless, recurring spectral spacings are used to start to unfold the rotational/tunneling structure. Moreover, spectral indicators are found that support the assumption of hydrazine-like tunnelling dynamics being present in this peculiar molecule of fundamental interest.

^a L. I. Yeh, M. Okumura, J. D. Myers, J. M. Price, and Y. T. Lee, Vibrational spectroscopy of the hydrated hydronium cluster ions $H_3O^+(H_2O)_n$ (n=1,2,3), J. Chem. Phys. 91, 7319-7330 (1989)

^b L. I. Yeh, Y. T. Lee, and J. T. Hougen. Vibration-rotation spectroscopy of the hydrated hydronium ions $H_5O_2^+$ and $H_9O_4^+$. Journal of Molecular Spectroscopy 164.2 (1994): 473-488.