Joint Annual Meeting of ÖPG and SPS 2021



Contribution ID: 12 Type: Poster

[631] Hyperfine effects in the vibrational spectroscopy of Cl⁻(H₂) and Cl⁻(D₂) complexes

Tuesday, 31 August 2021 19:01 (1 minute)

The predissociation spectra of the 35 Cl⁻(H₂) and 35 Cl⁻(D₂) complexes are measured at low frequencies between 400 and 800 cm⁻¹ in an ion trap at different temperatures. Above a certain temperature, the ligand switching between the two isotopologues ortho and para leads to a strong suppression of the excited hyperfine configuration. Performing the experiment below 30 K and 22 K for Cl⁻(H₂) and Cl⁻(D₂), respectively, we can detect the more weakly bound complexes.

Due to accurate quantum calculations, the bands in the $Cl^-(H_2)$ complex have been assigned to the intermonomer vibrational stretching mode.

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Session Classification: Poster Session

Track Classification: Applied Physics and Plasma Physics