

High-resolution anion momentum imaging study of the threshold dynamics of dissociative electron attachment to NO₂

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Dynamics near the molecular dissociation threshold can be complicated, if several close-lying electronic states or nuclear motions in multiple freedoms are involved. Experimental studies on this issue are frequently frustrated, due to low cross sections and demanding a high energy resolution to distinguish the manifold internal-energy distributions. Using the high-resolution anion velocity map imaging technique, here we report a dynamics study of the dissociations of anionic nitrogen dioxide in the low-lying resonant states formed by electron attachment [$e^- + NO_2 (X^2A_1) \rightarrow NO_2^- \rightarrow NO(X^2\Pi) + O^-(^2P)$]. The long-term puzzling issues about near-threshold dissociations of NO₂⁻ are settled in the present work. With help of theoretical computations, we suggest that three low-lying resonant states (¹B₁, ³B₁, ³B₂) of NO₂⁻ contribute to the O⁻ productions for the electron attachment energies less than 5 eV. Furthermore, B₁ and B₂ symmetries of the resonant states lead to different anisotropies of the O⁻ angular distribution. At the relatively high electron attachment energies, a prompt fragmentation in the molecular bent conformation competes with an indirect dissociation in the straightened conformation.

References:

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