

Resonant Raman scattering in 1s-shell photoionization of O₂ molecules

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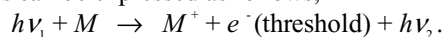
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Introduction

Doubly charged ions or more are observed in inner-shell photoionization of atoms. However, singly charged ions in 1s-shell photoionization of Ne have been observed by means of coincidence spectroscopy with ions and threshold photoelectrons [1]. Normally, Auger decays following inner-shell photoionization result in the formation of doubly charged ions or more. On other hand, radiative decays instead of Auger decays take part in the formation of singly charged ions. Namely, the decay process can be expressed as follows,



This process is the so-called continuous radiative resonant Raman scattering [2].

In inner-shell photoionization of N₂ molecules, singly charged ions, that is parent ions of molecules, have been also observed in coincidence with threshold photoelectrons [3]. The spectral structure was revealed by the radiative decays.

Experimental method

Experiments were carried out at the undulator beamline BL-2C. The beam line is equipped with a grazing incidence soft X-ray monochromator. A varied space plane grating with 1000 lines/mm was used. The spectral resolution of the monochromator with 50 μ × 10 μ slits was about 0.15 eV at 550 eV photon energy. Multiply charged ions in coincidence with threshold electrons were measured using a time-of-flight mass spectrometer coupled with a threshold-electron energy analyzer. The energy resolution of the analyzer was estimated to be about 0.03 eV.

Results and discussion

Figure 1 shows the yield spectra of ions, threshold photoelectrons and O₂⁺, O⁺ and (O⁺⁺ + O⁺⁺⁺) ions in coincidence with threshold photoelectrons near the 1s-shell photoionization region of O₂ molecules. The total yield spectrum of ions is correspond to the photoabsorption spectrum, and exhibits some resonance peaks of 1s-shell photoexcitation. The yield spectra of threshold photoelectrons and ions in coincidence with the threshold photoelectrons exhibit broad and distorted profiles due to post-collision interaction [4] in the 1s-shell photoionization threshold, in addition to some resonance peaks. However, the profile of parent ions O₂⁺ has symmetry just in the ionization threshold. Therefore, the

formation of parent ions is due to continuous radiative resonant Raman scattering [2].

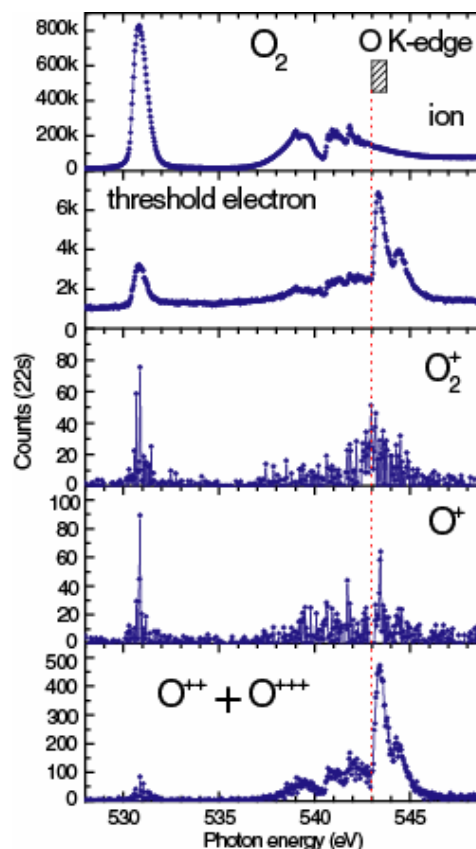


Figure 1. Yield spectra of ions, threshold electrons and ions in coincidence with threshold electrons near the 1s-shell photoionization region of O₂ molecules.

References

- [1] E. Murakami *et al.*, PF Activity Rep. **20**, B, 12 (2002).
- [2] T. Åberg and B. Crasemann, "Resonant anomalous X-ray scattering" pp. 431-448 (Elsevier Science) (1994).
- [3] T. Hayaishi *et al.*, PF Activity Rep. **22**, B, 2 (2004).
- [4] T. Hayaishi *et al.*, J. Phys. B. **27**, L115 (1994).

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