Resonant Auger Raman Spectroscopy of the Kr 3p->nl States

K.C. Prince¹, M. Coreno² and M. de Simone³,

¹Sincrotrone Trieste, Strada Statale 14, km. 163.5, in Area Science Park, I-34012 Basovizza (Trieste) ITALY

²INFM- TASC, Laboratorio ELETTRA, I-34012 Trieste, ITALY ³Università di Roma III ed Unità INFM, Via della Vasca Navale 84, I-00146 Rome, ITALY

X-ray absorption spectra of the sharp core levels of noble gases (Ne 1s, Ar 2p, Kr 3d, etc,) have been extensively studied and are well understood [1, 2,] but less attention has been given to the broader noble gas core levels [3, 4]. At the Kr 3p edge, the Rydberg sates are about 1.4 eV wide, overlap one another and are superimposed on a strong 3d continuum [4]. Thus the peaks are difficult to identify and controversy exists about their assignment [5].

The 3p⁻¹nl (n>4, l=0; n>3, l=2) Rydberg states decay rapidly via Coster-Kronig (CK) processes to 3d⁻¹4p⁻¹nl, 3d⁻¹4s⁻¹nl and related shake states, and via super Coster-Kronig processes. We have identified the CK states which lie close to the 3d⁻¹4p⁻¹np satellite states of the 3d line. On the first main 3p⁻¹nl resonance, there is a strong increase in emission below and above the 3d satellites. The peaks have widths determined by the resolution of the photons and the energy analysers, about 200 meV, fulfilling Resonant Auger Raman conditions. The new peaks disperse with photon energy as expected.

Constant Initial State spectra were measured by setting the analysers to energies corresponding to each group of peaks. The CIS spectra show clear differences in the resonant behaviour for different final state. One group of peaks show similar resonant behaviour on the absorption peak that has been attributed to the 3p⁻¹5s, both for 3p_{3/2} and 3p_{1/2} at 209.8 and 218 eV photon energy respectively. In contrast two other multiplet groups show completely different resonances, and are assigned to ionic states derived from different intermediate excited states.

By studying the decay states under Resonant Auger Raman conditions and using CIS spectroscopy, we have been able to separate the intermediate states in x-ray absorption at the 3p edge.

- [2] O.P. Sairanen et al, Phys. Rev. A **54** (1992) 2834.
- [3] T. Kylli et al, Phys. Rev. A. **59** (1999) 4071.
- [4] I.T. Steinberger et al, Phys. Rev. B **60** (1999) 3995.
- [5] M. Ohno, Phys. Rev. A **51** (1995) 1042.