Observation of interatomic Coulombic decay in Ne dimer above and below the Ne $2s^{-1}$ threshold

T. Aoto,^{*1} K. Ito,¹ Y. Hikosaka,² F. Penent,³ and P. Lablanquie,³ ¹KEK-PF, Tsukuba, Ibaraki 305-0801, Japan ² UVSOR Facility, Institute for Molecular Science, Okazaki 444-8585, Japan ³ LCP-MR, Université Pierre et Marie Curie, 11 rue P et M Curie, 75231 Paris, France

Introduction

It is well now that, in rare gas atoms, inner-shell core holes usually decay by emitting a secondary Auger electron. This pathway is forbidden in case of inner valence holes, for their binding energy lies below the lowest double ionization potential. Consequently, they can only experience a radiative decay. However, when the rare gas atom is a part of a weakly bound van der Waals cluster, an alternative interesting decay path is opened whereby the initially ionized atom transfers its energy to a neighboring atom. We have demonstrated that this process occur in Ar, Kr and Xe dimers [1]. The Ne dimer case is even more spectacular as the transferred energy is sufficient to ionize the neighboring atom. Such a decay process is called Inter-atomic Coulombic Decay (ICD) and was for the first time theoretically predicted by Cederbaum and co-workers [2,3] to be much faster than the competing intra-atomic radiative decay.

Results and Discussion

Ne dimer is the most suitable species in order to investigate the ICD mechanism. The process can be schematically depicted in the following way: an innervalence 2s electron is photoionized from a Ne atom inside the Ne dimer ([Ne---Ne]), the hole then decays by transferring its energy to the neighboring Ne atom and ejecting an outer-valence 2p electron, that is:

$$[\text{Ne---Ne}] + \text{hv}$$

$$\stackrel{\text{photoionization}}{\longrightarrow} [\text{Ne}^{+}(2s^{-1}) - --\text{Ne}] + e_{\text{photoelectron}}$$

$$\stackrel{\text{ICD}}{\longrightarrow} [\text{Ne}^{+}(2p^{-1}) - --\text{Ne}^{+}(2p^{-1})] + e_{\text{ICD}}.$$

The final state is thus a doubly charged Ne dimer that dissociates through charge separation. Recently, ICD was observed experimentally in 2s ionized Ne dimer by detecting the two Ne⁺ fragments and the ICD electron in coincidence at an excitation energy 10eV above the 2s ionization threshold [4].

In the present study, we used a very simple and sensitive experimental approach to investigate ICD process in Ne dimer, namely coincidence between the two receding energetic Ne⁺ ions. Experiment was

performed at undulator beamline BL16B. Ne dimers are produced in a supersonic expansion with a proper stagnation pressure in order not to produce large size clusters. The two fragment ions were detected in coincidence by two apparatus: a simple ion filter that rejects low energy or thermal ions and a hemispherical electrostatic analyzer [5] which allows the measurement of the kinetic energy distribution of energetic Ne^+ ions. We confirmed the ICD occurrence by detecting two Ne⁺ ions simultaneously with the two detectors. The ICD cross section were measured as a function of the photon energy, which is strongly affected by resonances due to double excitations, $[Ne^{(2p)^{4}(3s)(np)}]$, located just above the ionisation threshold of the $[Ne^{+}(2s^{-1})--Ne]$ state. The Ne⁺ kinetic energy observed in the present study is in good agreement with the previous theoretical prediction [3] and experimental observation [4] and remains constant within a few eV above threshold. Surprisingly, we observed that ICD process is also present below the 2s ionization threshold. Work is in progress to understand this phenomenon which we attribute to a spectator process upon excitation of a 2s electron to a Rydberg orbital.

References

[1] R. Thissen, P. Lablanquie, R. I. Hall, M. Ukai, and K. Ito, Eur. Phys. J. D 4, 335 (1998)

[2] L. S. Cederbaum, J. Zobeley, and F. Tarantelli, Phys. Rev. Lett. 79, 4778 (1997)

- [3] S. Scheit et al, J. Chem. Phys. 121,8393 (2004)
- [4] T. Jahnke et al, Phys. Rev. Lett. 93, 163401 (2004)
- [5] Y. Hikosaka, F. Penent, P. Lablanquie, R. Hall and K.

Ito, Meas. Sci. Technol. 11, 1697 (2000)

*tomohiro.aoto@kek.jp