Lawrence Berkeley National Laboratory

Recent Work

Title

ELECTRON IMPACT EXCITATION EFFICIENCY CURVES FOR THE FORMATION OF NEUTRAL METASTABLE SPECIES

Permalink https://escholarship.org/uc/item/7rk6s27z

Authors

Newton, Amos Thomas, G.E.

Publication Date

1970-06-01

For 18th Annual Conference on Mass Spectrometry and Allied Topics, San Francisco, Ca., June 14-19, 1970

LAWRENCE

LIBRARY AND DOCUMENTS SECTION UCRL-19594 Preprint

c.7.

RECEIVED ELECTRON IMPACT EXCITATION EFFICIENCY RADIATION LABORATORY CURVES FOR THE FORMATION OF NEUTRAL METASTABLE SPECIES JUL 29 1970

Amos S. Newton and G. E. Thomas

June 1970

AEC Contract No. W-7405-eng-48

TWO-WEEK LOAN COPY

This is a Library Circulating Copy which may be borrowed for two weeks. For a personal retention copy, call Tech. Info. Division, Ext. 5545

LAWRENCE RADIATION LABORATO NIVERSITY of CALIFORNIA BERKELEY

DISCLAIMER

This document was prepared as an account of work sponsored by the United States Government. While this document is believed to contain correct information, neither the United States Government nor any agency thereof, nor the Regents of the University of California, nor any of their employees, makes any warranty, express or implied, or assumes any legal responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by its trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof, or the Regents of the University of California. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof or the Regents of the University of California. ELECTRON IMPACT EXCITATION EMFICIENCY CURVES FOR THE FORMATION OF NEUTRAL METASTABLE SPECIES

Amos S. Newton and G. E. Thomas

Lawrence Radiation Laboratory University of California Berkeley, California 94720

For some years now, a group at the Lawrence Radiation Laboratory at Berkeley has been engaged in making measurements of the excitation efficiency curves for the formation of neutral electronically metastable species.^{1,2} Basically, an electron beam from an RPD gun has been crossed with an atomic or molecular beam. The metastable neutrals can be detected via their ability to eject electrons from a metal surface. In this work, the metal surface is the Cu-Be first dynode of an EMI type 9603 electron multiplier.

A difficulty encountered in making these measurements in this way is that U.V. photons produced in the collision chamber are also detected by the multiplier. However, the photons can be separated from the metastables by taking advantage of their flight time from the collision region to the detector. If excitation is effected by a short pulse of electrons, photons are observed to arrive at the detector during the pulse. The metastables arrive at later times.²

In attempting to extend the earlier work,^{1,2} we have developed a method of making RPD measurements on the time-resolved species. The pulsing scheme used is shown in Fig. 1. A master clock pulse triggers an 1 kHz pulse train from a time-based



Fig. 1. Pulsing scheme for time-resolved RPD measurements,

This work was performed under the auspices of the U.S. Atomic Energy Commission.

oscillator. These time-base oscillator pulses are used to trigger the channel advance of a Hewlett-Packard 5400 A multichannel analyzer used in the multiscale mode. An ~10 usec pulse, triggered by the time-base oscillator is applied to the RPD grid of a 5-electrode RPD gun. The voltage applied during the pulse is exactly that to allow a truncated electron distribution to pass through the collision chamber. The voltage of the collision chamber is set to give the desired initial electron energy. A data count gate is opened after a preset delay. The width and delay of the count gate is variable. This allows counts to be registered in the multichannel analyzer only during a specified time slot after the electron pulse. After VI msec, a second time-base oscillator pulse advances the channel number and advances the electron energy (derived from the X-output of the multi channel analyzer) by one step. After scanning 256 channels in this fashion, the multichannel analyzer produces an end-of-program pulse which stops the sequence. On receipt of the second clock pulse, the entire sequence is repeated, except that the electron 'on' voltage is lowered by VO.1 V, and the analyzer is automatically reset to operate in the subtract mode. This produces an RPD scan of the time-resolved species. Multiple RPD scans are always accumulated in the analyzer.

UCRL-19594





Fig. 2. Excitation efficiency curves for metastables and photons from N_2 .

curve in Fig. 2 shows the formation of metastables in N₂. This species has been examined and discussed in detail by, e.g., Olmsted,³ Lichten,⁴ and Freund.⁵ In the present study, the $E^{3}\Sigma_{g}^{+}$ state near 12 eV shows up clearly. The lower curve of Fig. 2 shows the photon signal observed in N₂. The appearance potential of the photons coupled with the knowledge that they are sufficiently energetic to eject an electron from the multiplier strongly suggests that the Birge-Hopfield System is the source of the light.

-3-

In the studies of the rare gases Ne, Ar, and Kr, the metastable excitation efficiency curves agree very well with those of Olmsted, Newton, and Street.¹ In all cases, the photon appearance potential was equal, within experimental error, to that of the metastable. In Ar and Kr, a break in the photon curve was observed near the energy of the $np^{5}nd$ configurations.

A time-of-flight survey of a variety of gases has revealed that there are species with a flight time shorter than that expected for the molecular metastable (but with a flight time longer than the fast photon signal) in 0_2 and in CS_2 . No excitation efficiency curves for the latter species are available.

References

John Olmsted III, Amos S. Newton, and K. Street, Jr., J. Chem. Phys. <u>42</u>, 2321 (1965).
R. Clampitt and Amos S. Newton, J. Chem. Phys. <u>50</u>, 1997 (1969).

3. J. Olmsted III, Radiation Res. <u>31</u>, 191 (1967).

4. William Lichten, J. Chem. Phys. 26, 306 (1957).

5. Robert S. Freund, J. Chem. Phys. 50, 3734 (1969).

LEGAL NOTICE

This report was prepared as an account of Government sponsored work. Neither the United States, nor the Commission, nor any person acting on behalf of the Commission:

- A. Makes any warranty or representation, expressed or implied, with respect to the accuracy, completeness, or usefulness of the information contained in this report, or that the use of any information, apparatus, method, or process disclosed in this report may not infringe privately owned rights; or
- B. Assumes any liabilities with respect to the use of, or for damages resulting from the use of any information, apparatus, method, or process disclosed in this report.

As used in the above, "person acting on behalf of the Commission" includes any employee or contractor of the Commission, or employee of such contractor, to the extent that such employee or contractor of the Commission, or employee of such contractor prepares, disseminates, or provides access to, any information pursuant to his employment or contract with the Commission, or his employment with such contractor. TECHNICAL INFORMATION DIVISION LAWRENCE RADIATION LABORATORY UNIVERSITY OF CALIFORNIA BERKELEY, CALIFORNIA 94720

N)