

A Cryogenic Cylindrical Ion Trap Velocity Map Imaging

Spectrometer

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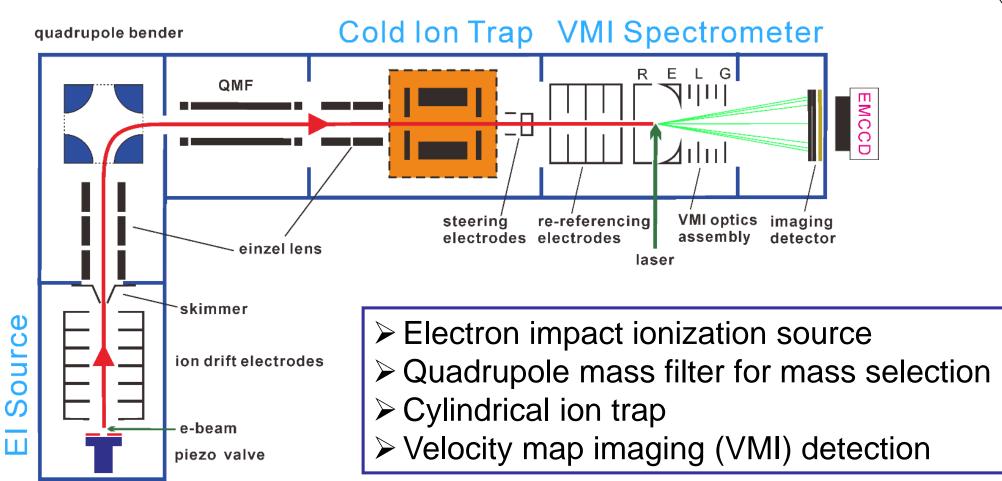
A cryogenic cylindrical ion trap velocity map imaging spectrometer (CIT-VMI) has been developed to study photodissociation spectroscopy and dynamics of gaseous molecular ions and ionic complexes. A cylindrical ion trap (CIT) made of oxygen-free copper is cryogenically cooled down to ~7 K by a closed cycle helium refrigerator and is coupled to a velocity map imaging (VMI) spectrometer. For CO₂+ ions, a rotational temperature of ~12 K is estimated from the recorded [1+1] two-photon dissociation spectrum, and populations in spin-orbit excited X²Π_{α,1/2} and vibrationally excited states of CO₂+ are found to be non-detectable, indicating an efficient internal cooling of the trapped ions. Based on the time-of-flight peak profile and the image of N_3 , the velocity spread of the ions extracted from the trap, both radially and axially, is interpreted as approximately ± 25 m/s. Experimental image of fragmented Ar+ from 307 nm photodissociation of Ar₂+ shows that, benefitting from the well-confined velocity spread of the cold Ar₂+ ions, a VMI resolution of Δv/v ~ 2.2% has been obtained.

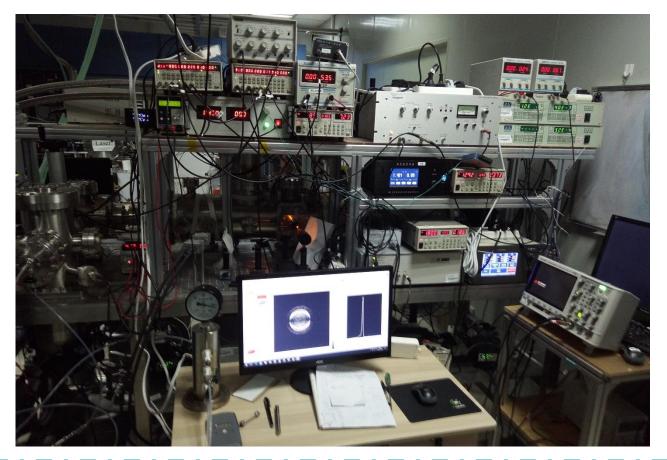
Introduction

A combination of radio frequency cylindrical ion trap and velocity map imaging make it possible to study high resolution photodissociation spectroscopy and dynamics of molecular and cluster ions by an ion beam with significantly reduced speed spread.

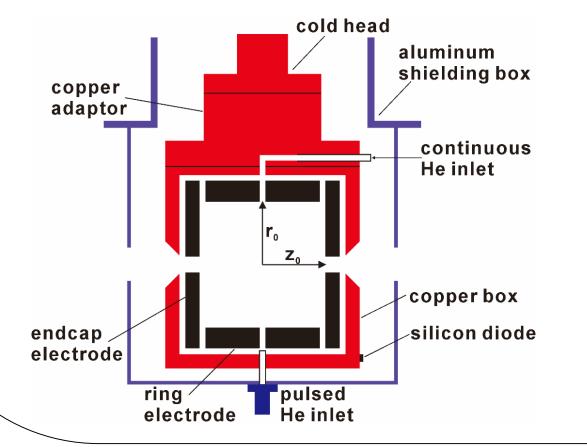
- Radio frequency cylindrical ion trap with cryogenic cooling
- Velocity map imaging

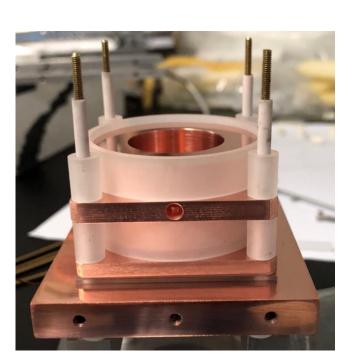
Experimental setup



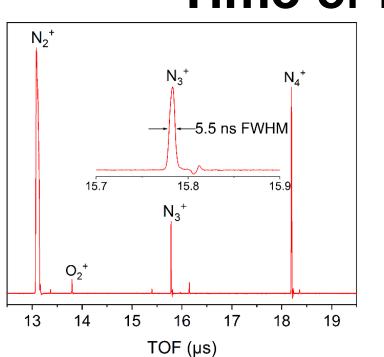


Cylindrical ion trap (CIT)





Time-of-flight mass spectra



electron ionization of N_2 .

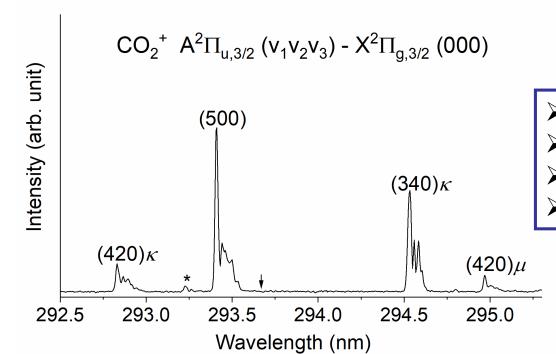
TOF mass spectra of N_x^+ (x=2-4) by

The observed mass peaks in the TOF spectra.

| m/z | Time (μ s) | FWHM (ns) | Resolution $(m/\Delta m)$ |
|-----|-----------------|-----------|---------------------------|
| 28 | 13.10 | 48.7 | 134 |
| 32 | 13.80 | 5.9 | 1169 |
| 42 | 15.78 | 5.5 | 1435 |
| 44 | 16.15 | 6.2 | 1302 |
| 56 | 18.20 | 9.9 | 919 |

➤ Initial axial size: ~0.3 mm ➤ Axial speed spread: ± 25 m/s

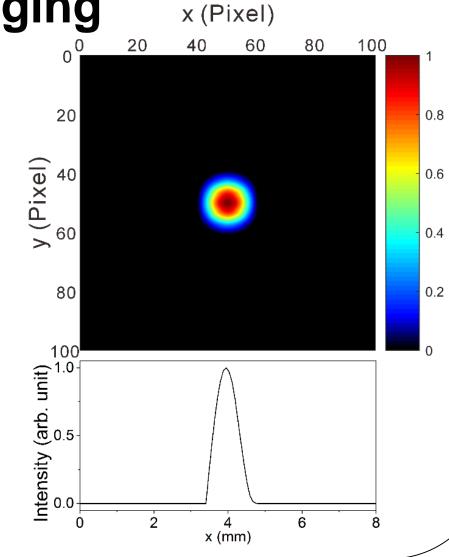
Photodissociation spectroscopy



- > Rotational temperature: 12 ± 2 K ➤ No detection of electronic excited CO₂+ ➤ No detection of vibrational excited CO₂+
- \triangleright No detection of CO₂+ X² $\Pi_{a_1/2}$ (000)
- [1+1] two-photon dissociation spectrum of CO₂+.

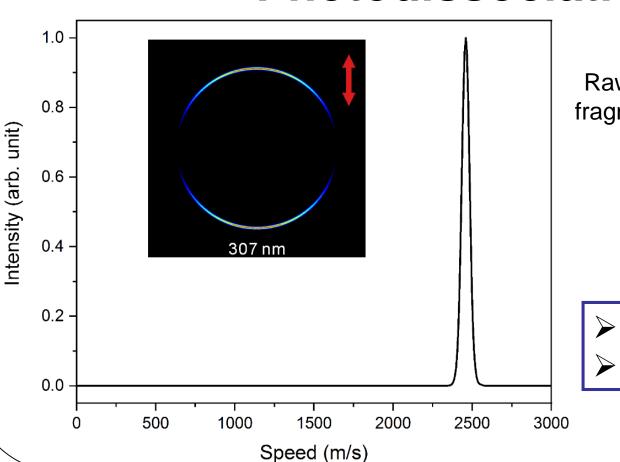
Ion imaging

➤ Initial radial size: ~0.2 mm ➤ Radial speed spread: ± 25 m/s



Upper panel: the image of N_3^+ ions after extraction from the trap; lower panel: the distribution of N_3^+ ions along the x axis.

Photodissociation of Ar₂+



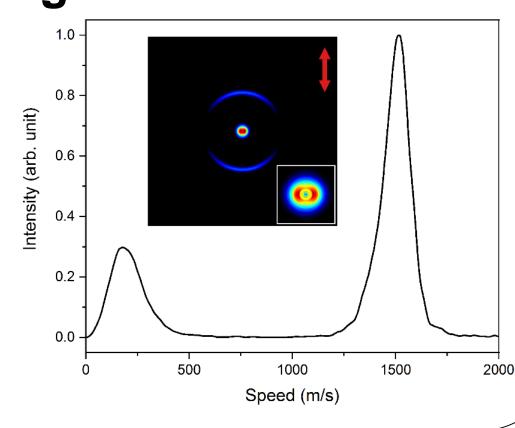
Raw image and speed distribution of fragmented Ar+ from photodissociation of Ar_2^+ at 307 nm.

 \triangleright VMI resolution: $\Delta v/v \sim 2.2\%$ \geqslant β =1.9 , parallel transition

Work in progress

 $Ar_3^+ + hv$ \rightarrow 2Ar + Ar+

Raw image and speed distribution of fragmented Ar+ from photodissociation of Ar_3^+ at 529 nm.



References

Z. Hua, S. Feng, Z. Zhou, H. Liang, Y. Chen, D. Zhao, Rev. Sci. Instrum. 2019, 90, 013101.

A. T. J. B. Eppink, and D. H. Parker, Rev. Sci. Instrum. 1997, 68, 3477.

C. R. Gebhardt, T. P. Rakitzis, P. C. Samartzis, V. Ladopoulos, T. N. Kitsopoulos, Rev. Sci. Instrum. 2001, 72, 3848. M. A. Gharaibeh and D. J. Clouthier, J. Chem. Phys. 2010, 132, 114307.

Acknowledgements

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