

# Highly Efficient TOF-Measurements of Photoelectrons and -ions at the PETRA III-XUV Beamline

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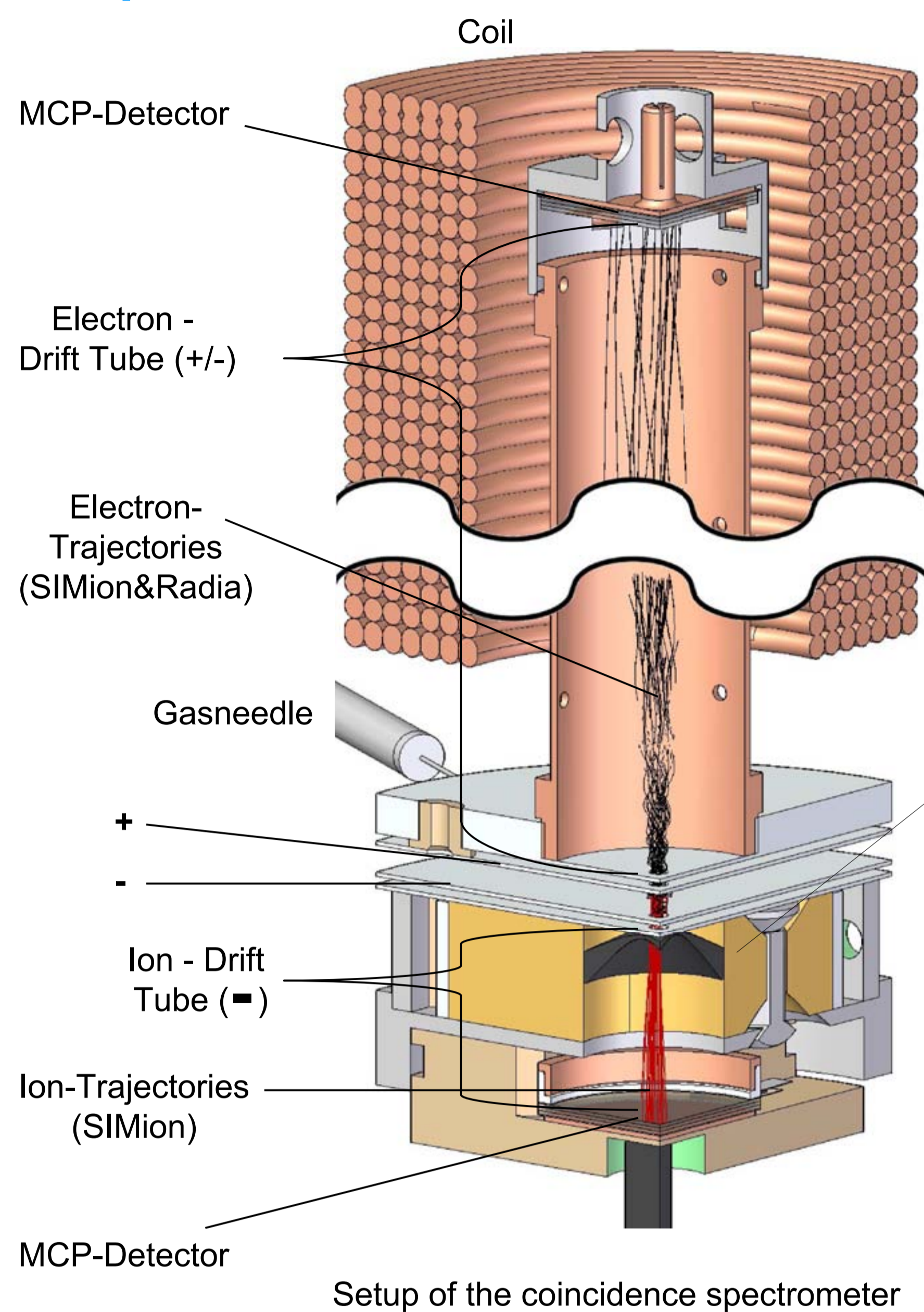
## Introduction

A highly efficient Time-of-Flight spectrometer for coincident photoelectron and -ion measurements has been used for various photoionization experiments at the P04 Variable Polarization XUV Beamline (PETRA III). The spectrometer consists of a short Time-of-Flight spectrometer for ion detection, using electrostatic fields according to the principle of Wiley and McLaren [1], and a so called *magnetic bottle* spectrometer for efficient electron measurement [2]. The design is based on the work of Eland and Feifel [3] but it is optimized for the specifications of the P04-Beamline.

## Ion-ToF-Spectroscopy

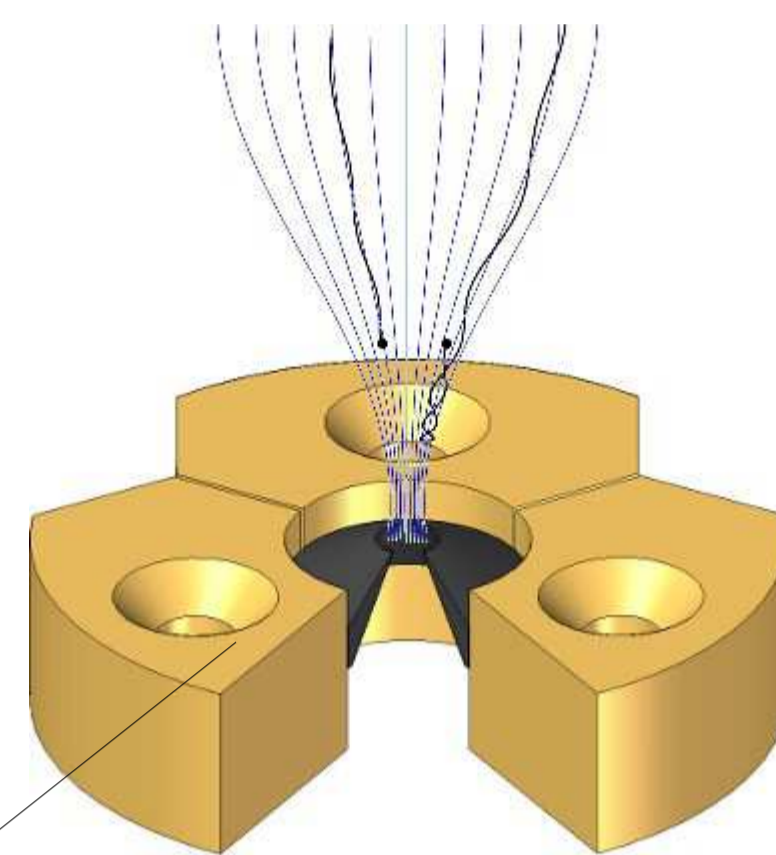
Various ion-ToF-spectra of atoms (Ne, Ar, Xe...) and molecules (H<sub>2</sub>, SF<sub>6</sub>, CO<sub>2</sub>...) have been measured, e.g. see poster 39 (Gregor Hartmann et al.).

## Setup



## Characteristics

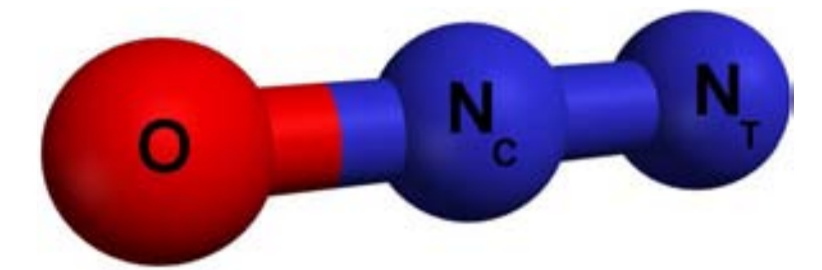
- Energy Resolution 1-2 eV
- Mass Resolution ~250
- Efficiency (Electrons) up to 30 %
- Efficiency (Ions) up to 55 %



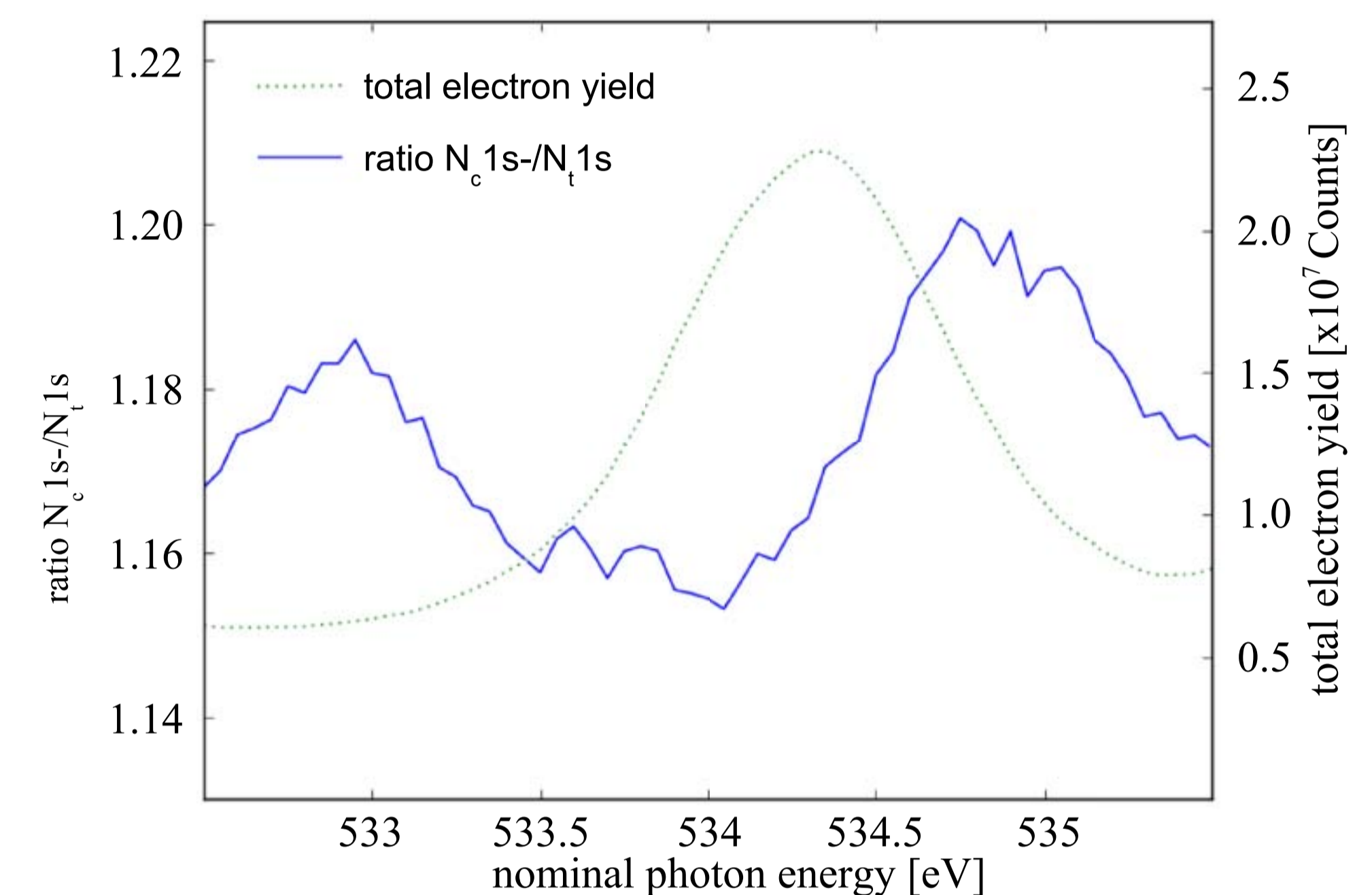
Assembly of three magnets and pole cap generating the magnetic field. The bottle neck shape turns electron trajectories upwards to the detector while the kinetic energy is conserved.

## Electron-ToF-Spectroscopy on N<sub>2</sub>O

### NACHT – Effect<sup>[4]</sup>

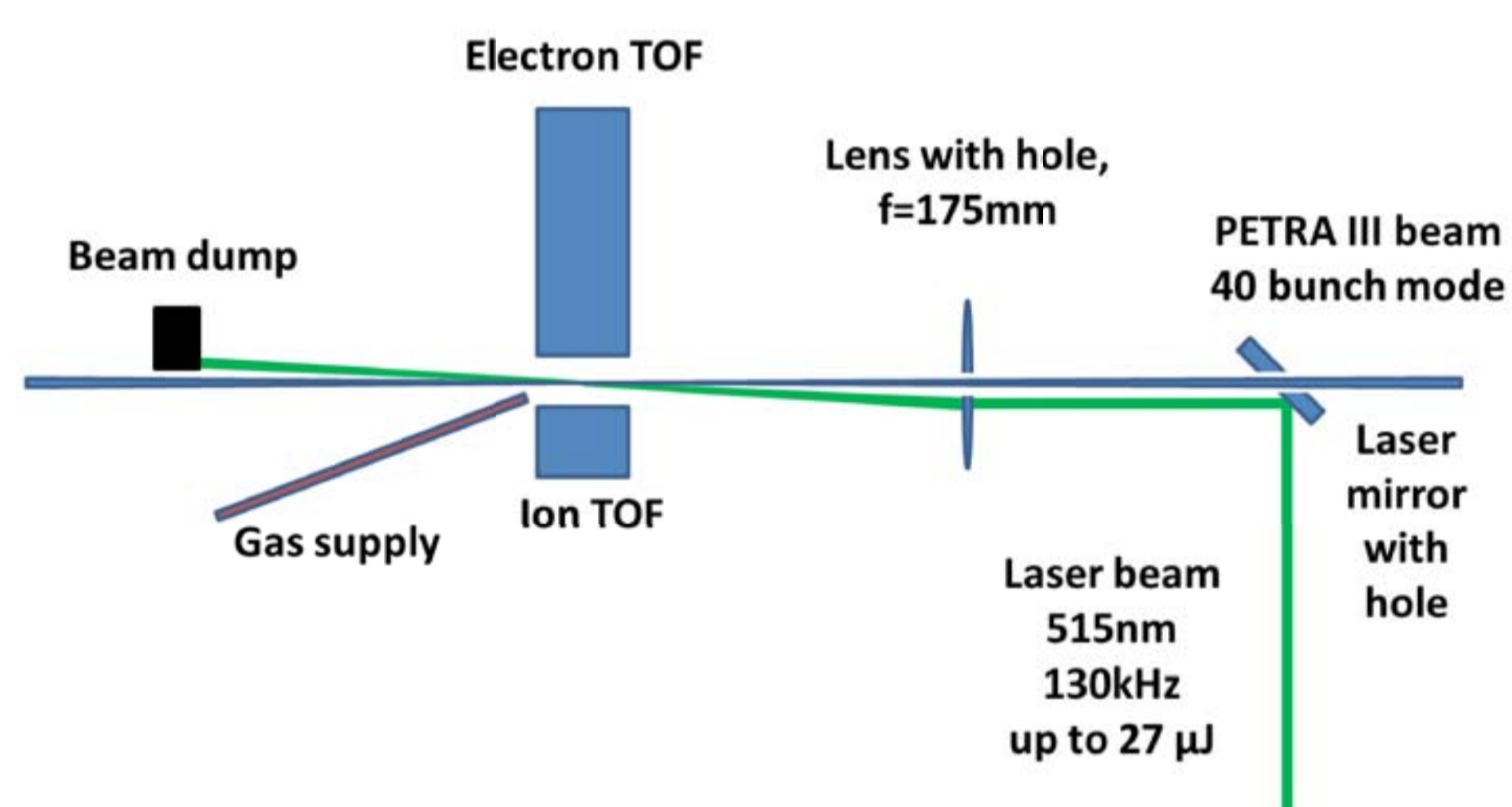


The chemical shift in N<sub>2</sub>O allows to distinguish 1s-photoelectrons from the central (E<sub>B</sub>(N<sub>c</sub>1s) = 412.5 eV) and the terminal nitrogen atom (E<sub>B</sub>(N<sub>t</sub>1s) = 408.5 eV). It has been shown that the O1s → 3π\* excitation leads to a change in the non-dipole parameter for the N1s-photoelectron emission [4]. The new revision of the magnetic bottle allows clear separation of the N<sub>c</sub>1s- and N<sub>t</sub>1s-photoelectrons in the range of the O1s → 3π\* resonance.

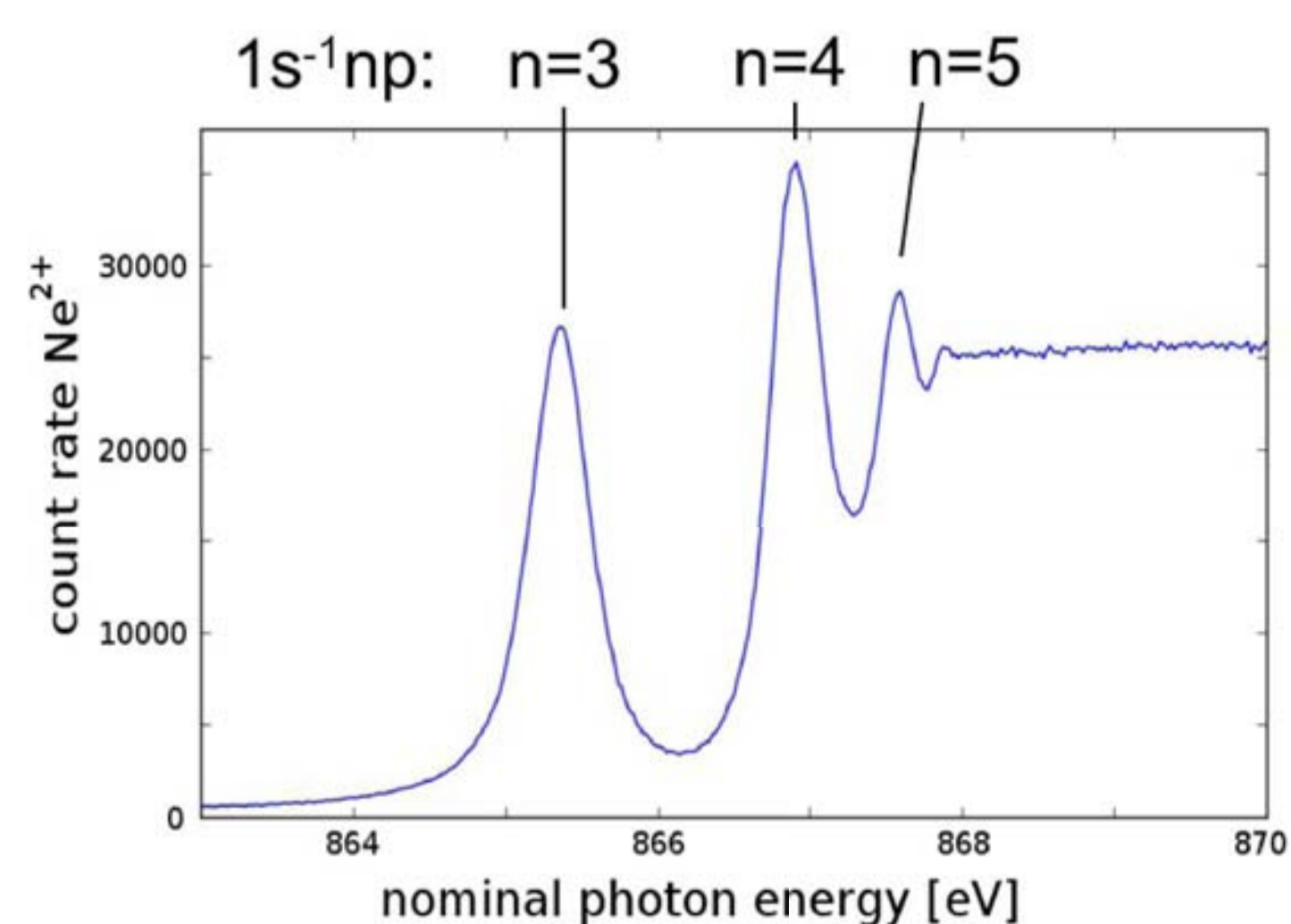


Total electron yield and ratio of N<sub>c</sub>1s- and N<sub>t</sub>1s- electrons in the range of the O1s → 3π\* resonance. The ratio changes with a shift to the resonance due to an increased N<sub>c</sub>1s- electron yield.

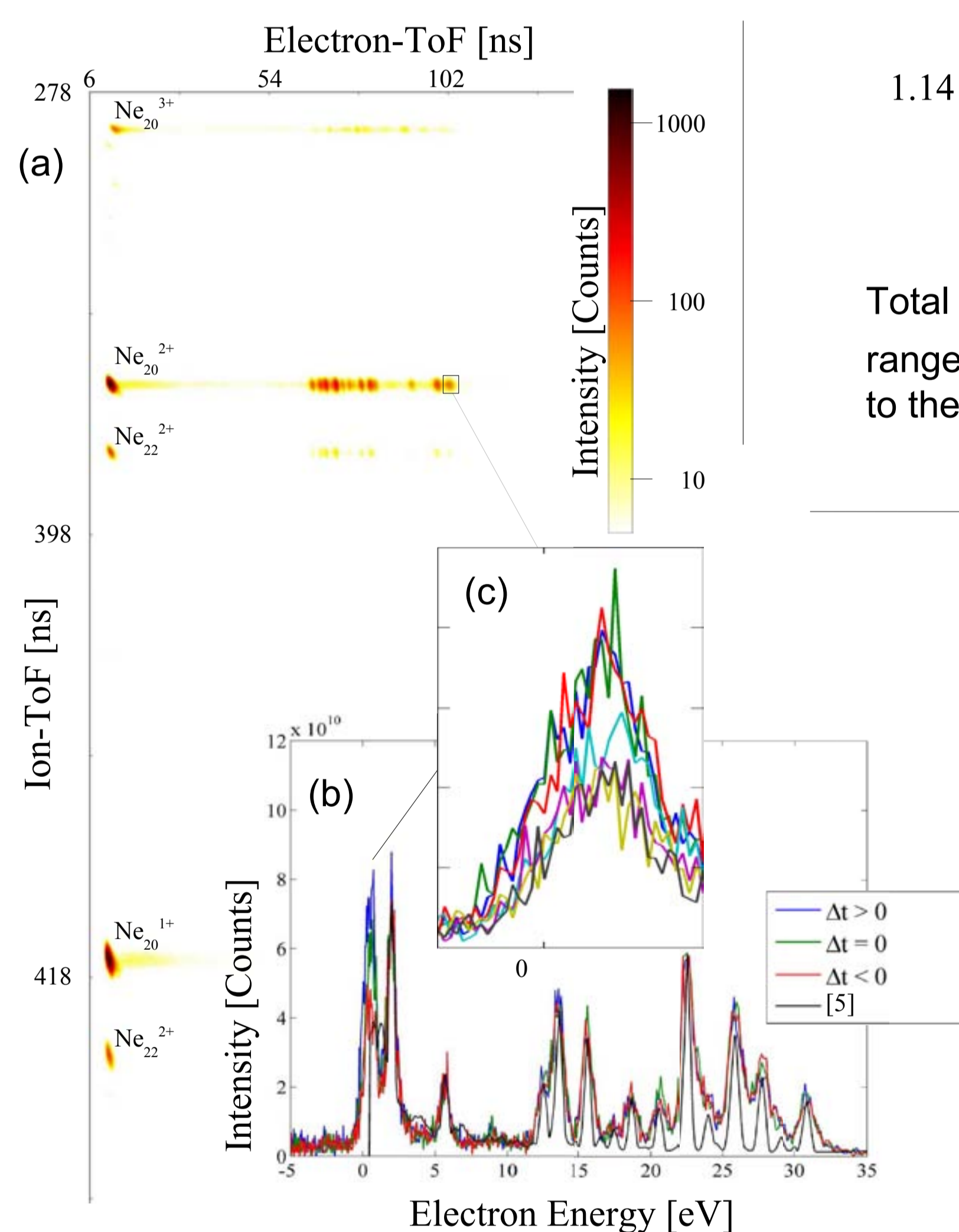
## Time Resolved Studies



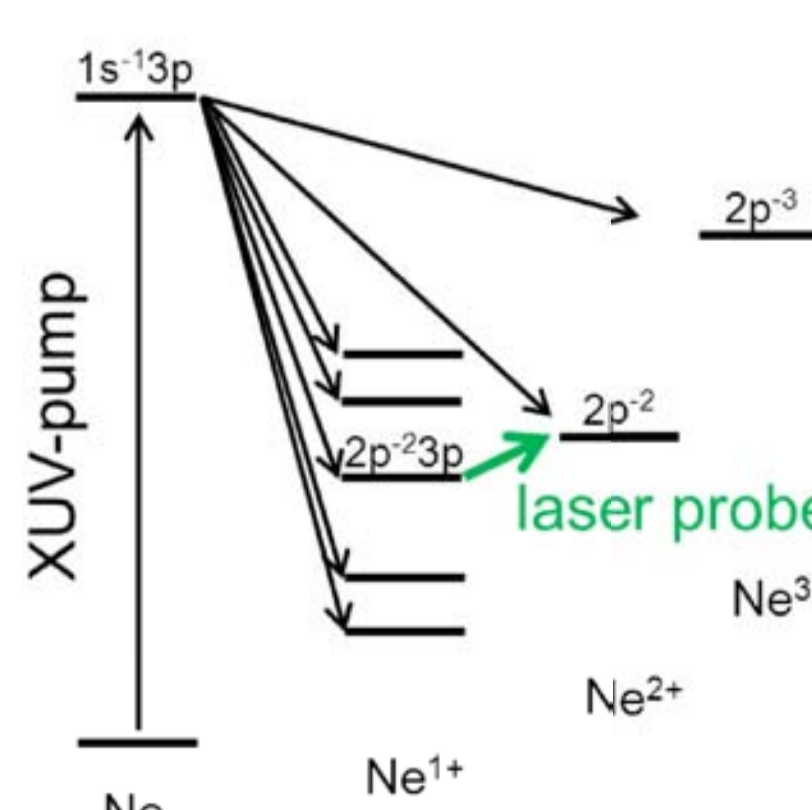
Schematic of the XUV-pump – laser probe experiment. The laser system is synchronized to the storage ring with a residual jitter of ~15ps. Every 40<sup>th</sup> bunch can be pumped at ~130kHz repetition rate.



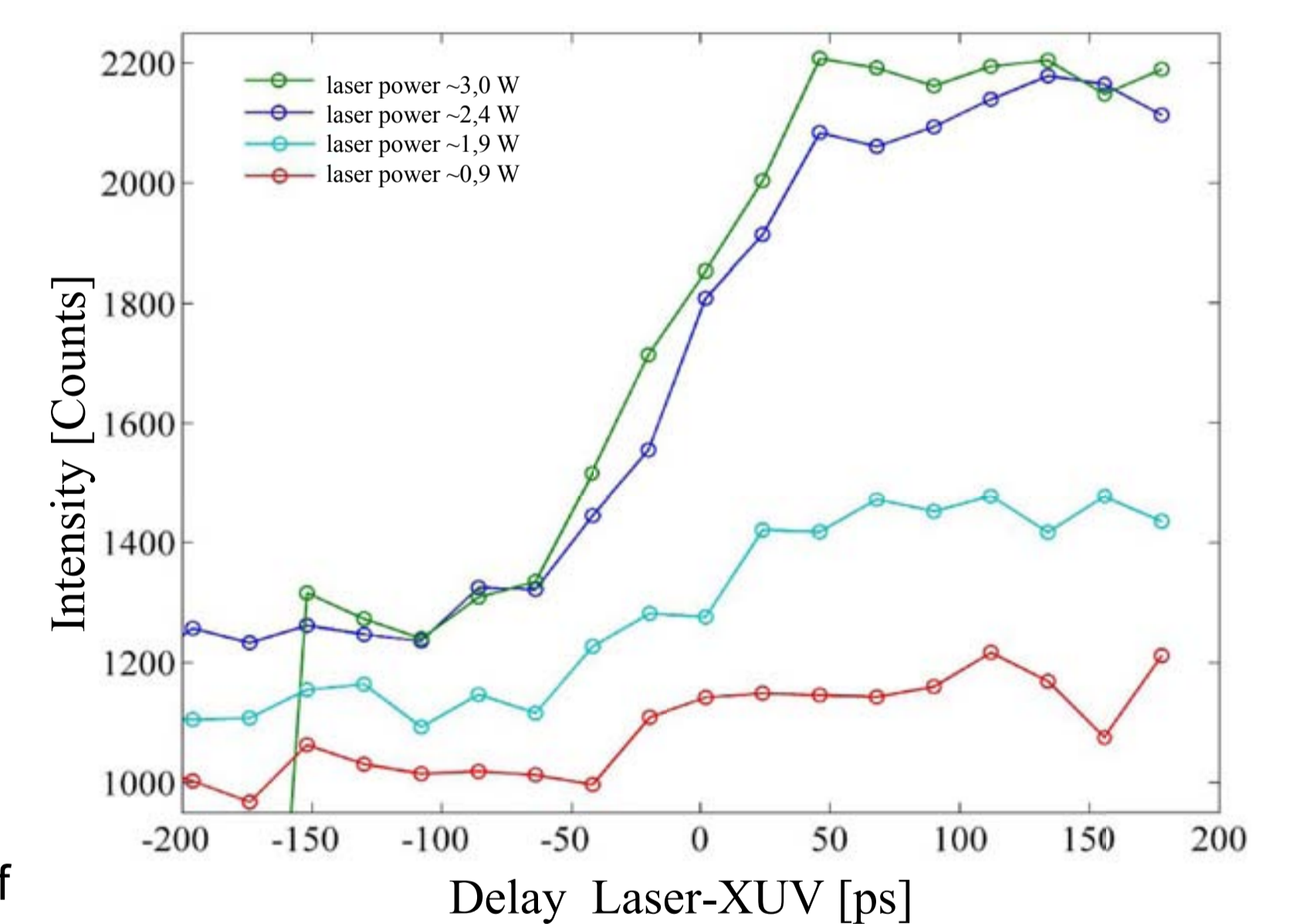
Ne<sup>2+</sup> ion yield at the Ne1s ionization threshold showing the first resonant rydberg states of the 1s-np-excitation.



Electron-ion coincidence map of Neon with a laser delay of Δt ~ 200 ps (a) and spectra of electrons in coincidence with Ne<sup>2+</sup> with different delays (b) compared with [5]. In (c) the slow electron signal is enlarged for different delays. All spectra were taken on the Ne 1s → 3p resonance.



Simplified energy level scheme of Ne. The laser-probed state is indicated by the green arrow.



Ion yield of Ne<sup>2+</sup> in coincidence with slow electrons vs. laser delay and laser power. The rise time of the signal corresponds to the pulse duration of the X-ray pulses of ~100 ps.

## References

- [1] P. Kruit, F.H. Read, J. Phys. E 16 (1983) 313
- [2] W.C. Wiley, I.H. McLaren, Rev. Sci. Instrum. 26 (1955) 1150
- [3] J.H.D. Eland, R. Feifel, Chemical Physics 327 (2006) 85–90
- [4] R. Guillemin et al., Phys. Rev. Lett. 92:223002 (2004)
- [5] J. Viefhaus, PhD Thesis