

EXPLORING THE RESOLUTION LIMIT OF PFI-ZEKE PHOTOELECTRON SPECTROSCOPY

H. HERBURGER, V. WIRTH U. HOLLENSTEIN, F. MERKT, *Department of Chemistry and Applied Biosciences, ETH Zurich, 8093 Zurich, Switzerland*

The spectral resolution in pulsed-field-ionisation zero-kinetic-energy (PFI-ZEKE) photoelectron spectroscopy is related to the state selectivity in the diabatic ionisation of the Rydberg states by time-dependent electric fields. The selectivity is determined by the applied electric-field pulse sequence. Hollenstein *et al.*¹ used discrete electric-field pulses with increasing field strength in combination with a preceding field pulse of opposite polarity and achieved a spectral resolution of 0.06 cm^{-1} . To improve the resolution further, Harper *et al.*² recently replaced the sequence of field steps by a linearly increasing field, as used earlier by Reiser *et al.*³, in combination with a prepulse of opposite polarity and obtained promising results on the PFI-ZEKE photoelectron spectrum of NO and CO₂. They called this new method pulsed-ramped field-ionisation (PRFI)-ZEKE photoelectron spectroscopy.

Using a home-built narrow-bandwidth long-pulse laser system (pulse lengths up to 50 ns, bandwidth < 50 MHz) in combination with a pulsed field ramp,^{2,3} we explore the resolution limit of this approach. To avoid overlap of spectral lines, we chose the atoms Ar, Kr, and Xe as test systems and recorded PRFI-ZEKE photoelectron spectra of transitions from the metastable state $(np)^5((n+1)s)[3/2]_2$ (³P₂) of the neutral rare-gas atom to the $(np)^5$ ²P_{3/2} state of the corresponding ion, with $n = 3, 4, 5$ for Ar, Kr, and Xe, respectively. We present PRFI-ZEKE photoelectron spectra that exhibit both adiabatic and diabatic ionisation dynamics. The resolution of these spectra is improved to 0.05 cm^{-1} which constitutes the resolution record in photoelectron spectroscopy.¹ The determined ionisation energies are compared with precisely known literature values⁴. The current resolution is limited by adiabatic field ionisation below $n = 190$ and by electric stray fields, which affect the field-ionisation dynamics above $n = 190$.

¹U. Hollenstein, R. Seiler, H. Schmutz, M. Andrist, and F. Merkt, *J. Chem. Phys.* **115**, 5461–5469 (2001).

²Oliver J. Harper, Ning L. Chen, Séverine Boyé-Péronne, and Bérenger Gans, *Phys. Chem. Chem. Phys.* **24**, 2777–2784 (2022).

³G. Reiser, W. Habenicht, K. Müller-Dethlefs, and E. W. Schlag, *Chem. Phys. Lett.* **152**, 119–123 (1988).

⁴V. L. Sukhorukov, I. D. Petrov, M. Schäfer, F. Merkt, M.-W. Ruf, and H. Hotop, *J. Phys. B: At. Mol. Opt. Phys.* **45**, 092001 (2012) and references therein.