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Time-resolved and XUV spectroscopy of helium nanodroplets

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Retraction

Time-resolved and XUV spectroscopy of helium nanodroplets

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Time-resolved and XUV spectroscopy of helium nanodroplets

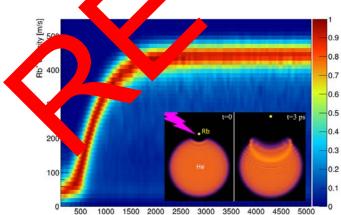
M. Mudrich¹*, J. von Vangerow¹, A. LaForge¹, F. Stienkemeier¹, A. Ciavardini², P. O'Keeffe², Y. Ovcharenko³, T. Möller³, M. Drabbels⁴, P. Piseri⁵, O. Plekan⁶, P. Finetti⁶, M. Coreno⁷, C. Grazioli⁶, R. Richter⁶, K. C. Prince⁶, C. Callegari⁶

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Synopsis The ultrafast dynamics of pure and doped helium nanodroplets is studied using VIS at XUV femtosecond pump-probe spectroscopy in combination with ion and electron imaging detection

Helium (He) nanodroplets currently attract considerable interest mainly for two reasons. 1) Due to their quantum fluid nature He nanodroplets feature extraordinary properties such as microscopic superfluidity. 2) He nanodroplets can be considered as a nearly ideal matrix for spectroscopy of embedded molecules and aggregates due to their transparency for light up to the extreme ultraviolet (XUV) spectral range, and their ability to efficiently aggregate and cool embedded species.

However, the short-time dynamics of He nanodroplets is still largely unexplored. Recently, strong perturbations of the rotational mo of embedded molecules induced by the droplet environment was observed for the ca of impulsive excitation [1]. In this ribution we study the ultrafast response o lase excitation of a prototype system consting of a lkali metal atom residing at the surf nanodroplets. Using fer tosecol oump-probe spectroscopy in combination with N and electron imaging deterion follow in val-time the desorption of the exclusive atom off the He droplet surface (see Fig. 1) a well as the formation of *2* ali-He exciplex more cules.



Pump-probe delay [fs]

Figure 1. Time-evolution of the velocity of an excited Rb atom ejected off the surface of a He nanodroplet. Inset: TDDFT simulation [2].

Upon irradiation of He *p*plets with at $hv \ge 21$ eV, where He drore its are satisfy absorbing, a complex poto mamics is initiated by the excitetion or zation of Ae atoms ets [3]. The lynamic of relaxainside the dr tion, fragmenta, n, as well an indirect ioniza-tion processes are indied using synchrotron and free-electron-laser (Pomi@Elettra) radiation. In cular, the real-time dynamics of pure He pa odroplets excited by tunable femtosecond n V-pulses is udied in a pump-probe exper-Σ t. The time resolved photoelectron spectra in revel ultrafation intra- as well as interband relaxation dynamics, see Fig. 2.

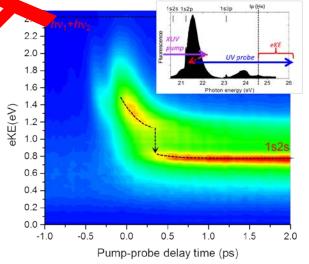


Figure 2. Pump-probe delay dependent photoelectron spectrum of pure He nanodroplets measured by XUV (22.2eV)/UV (4.8eV) fs pump-probe R2PI.

References

[1] D. Pentlehner et al., Phys. Rev. Lett. 110, 093002 (2013) [2] J. von Vangerow et al., J. Phys. Chem. A 118, 6604 (2014) [3] M. Mudrich and F. Stienkemeier, Int. Rev. Phys. Chem. 33, 301 (2014)

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