Few-photon multiple ionization of Ne and Ar by strong FLASH pulses

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Few-photon multiple ionization of Ne and Ar atoms by strong VUV laser pulses from the Free electron LASer at Hamburg (FLASH) was investigated differentially with the Heidelberg reaction microscope. The light intensity dependence of Ne²⁺ production reveals the dominance of non-sequential two-photon double ionization at intensities of $I < 6 \times 10^{12}$ W/cm² and significant contributions of sequential ionization as I increases. Ne²⁺ recoil-ion momentum distributions suggest that two electrons absorbing "instantaneously" two photons are ejected most likely into opposite hemispheres with similar energies.

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Single-photon induced multiple ionization or excitation of atoms and molecules has been central to atomic and molecular physics research due to their prototype character to explore electron-electron correlation. Experimental milestones were, among others, kinematically complete measurements on double ionization of He and H_2 (see e.g. [1, 2]), the fundamental two-electron systems. Here, quantum mechanical ab initio calculations have emerged since 1998, with predictions being in excellent agreement with all available data such that single-photon double-ionization of He is considered to be understood [3] within the validity of the dipole approximation. Multi-photon induced multiple ionization at the other extreme, where a couple of ten photons are typically needed using lasers in the visible, has until the present day resisted any comprehensive theoretical description due to tremendous complications in solving the non-perturbative, few-particle quantum problem (see e.g. [4]).

Few-photon multiple ionization, e.g. the interaction of two or three photons with two or more electrons, represents one of the most fundamental non-linear processes, bridges the gap between the single- and multi-photon regimes and, thus, is of decisive importance to advance non-linear theories. Mainly due to its perturbative nature at shorter wavelengths along with the fact that only few photons are involved, has allowed for the recent emergence of full quantum calculations [5].

The Free electron LASer at Hamburg (FLASH) [6] delivering vacuum ultraviolet (VUV) photons at unprecedented intensities, in combination with the most advanced multiparticle detection systems such as the Heidelberg reaction microscope [7], open the door for performing kinematically complete experiments in this new regime. At not too high intensities ($I \approx 10^{13}$ W/cm²), multi-photon ionization (MPI) can be expected to be the dominant process for many-electron removal. MPI contains two basic dynamical processes, sequential ionization (SI) and non-sequential ionization (NSI). NSI requires that electrons are ionized by simultaneous ab-

sorption of photons, whereas SI is a sequential absorption process within the same light pulse from FLASH. Since for SI, the intermediate ionic state is usually assumed to be relaxed in its electronic ground state before further photons are absorbed, typically more photons are needed than for NSI to reach the same final charge state. NSI on the other hand, due to its simultaneous rather than "step-by-step" character, provides a much richer setting for photon-electron interaction as well as electron-electron correlation mechanisms making it the more exciting process.

Differential experimental data for MPI represent the most critical testing ground for theories. Besides the basic character of such experiments, there is tremendous practical interest since these non-linear processes govern the interaction of the VUV-radiation with matter in general and have to be understood if more complicated reactions with surfaces, bulkmatter or biological samples shall be investigated.

Few experiments on total cross sections have been reported so far. Recently, two-photon single ionization of He was investigated by Laarmann *et al.* [8] with FLASH radiation at 98 nm wavelength. Nabekawa *et al.* [9] observed the production of doubly charged He ions by two-photon absorption using 42 eV high-harmonic radiation. Again at 98 nm with FLASH, Wabnitz *et al.* [10] studied multiple ionization of Ar and Xe atoms and experimentally determined multiply charged Xe^{*q*+} yields were interpreted in terms of the absorption of several VUV photons with the help of an independent-particle model [11]. Here, *q* denotes the charge-state of the ion. Very recently, Benis *et al.* [12] observed two-photon double ionization of Ar and Kr atoms by a superposition of harmonics.

In this letter, we present momentum-resolved experimental spectra for two-photon double ionization of Ne obtained with a reaction microscope at FLASH. The data provide first information about the sum-momentum of two emitted electrons supporting dominant NSI contributions at light intensities of a few 10^{12} W/cm² in accordance with the recorded intensity

dependence of Ne^{2+} ion-yields. With increasing intensity SI was found to become more and more important.

Our reaction microscope was installed at the focus of beam line BL 2 at FLASH. The light at 32 nm (\approx 38.8 eV), with a 5 Hz repetition rate at pulse energies \approx 1–10 μ J, was focused into the interaction chamber reaching peak intensities of $I \approx 10^{12} \sim 10^{13}$ W/cm². The FEL-beam was crossed with a well collimated (2 mm diameter), dilute (\approx 10⁹ particles/cm³) and intrinsically cold (T_{jet} \approx 2 K) supersonic atomic gas jet. Ions produced by the interaction with the FEL-pulse were projected by means of a weak electric field (0.24 V/cm) onto time- and position-sensitive MCP detectors (diameter 120 mm, position resolution 0.1 mm). From the measured timeof-flight and position of each individual particle the initial 3-dimensional momentum vectors were reconstructed and an ion-momentum resolution of \approx 0.8 a.u. for Ne⁺ was achieved (for details, see [7]).

The current induced on a metal plate by each individual FLASH-pulse was recorded in order to obtain pulse-to-pulse information of the respective pulse energy. By comparing the average induced current over several thousand pulses with the mean pulse energy given from the machine the absolute energy of each pulse could be deduced. On the basis of this number, the approximately known focal spot size (30 μ m diameter) and the pulse duration (30 fs), *I* was calculated with an estimated uncertainty of a factor of 5. The linearity of our intensity scale was assured to be correct within 10% with a relative calibration between runs of better than 50% by comparing different yields for single ionization (²⁰Ne⁺, ²²Ne⁺, H₂O⁺, etc.) as a function of *I*.

The time-of-flight spectra of Ne^{*q*+} for q = 1 and 2, and Ar^{*q*+} for q = 1-3, are shown in Fig. 1. The spectrum is dominated by peaks of singly charged ²⁰Ne⁺ and ²²Ne⁺ ions with small (~2%) admixtures from the residual ion H₂O⁺ (broad hump). In addition, small but significant yields of ²⁰Ne²⁺ ions were observed. Similarly to Ne, MPI of Ar producing up to Ar³⁺ was found (Fig. 1 inset).

In analyzing the data, one has to first clarify whether multiply charged Ne^{*q*+} and Ar^{*q*+} ions result from MPI or singlephoton absorption of high-harmonic radiation, which was not actively suppressed in the present experiment. On the basis of the well-known cross sections for single-photon single and double ionization of Ne by SR [13, 14], one can estimate the ionization-rate ratio $P_{\text{Ne}^{2+}}/P_{\text{Ne}^+}$ to be $\approx 7.5 \times 10^{-5}$ for single photon absorption from the second and third harmonics. For this estimate, the relative intensity ratios of the second and third harmonics to the fundamental were taken from Ref. [15] in the 10 – 40 nm range to be 2nd/1st=0.35% and 3rd/1st=0.4%. This result is in contrast to the measured ratios of 2.6×10^{-4} to about 2×10^{-3} providing evidence that the observed multi-charged ions mainly result from few-photon ionization.

The first and second ionization potentials for Ne are 21.6 eV and 40.9 eV respectively, and the first four ionization potentials of Ar are 15.8 eV, 27.7 eV, 40.7 eV, and 59.8 eV. Therefore, a minimum energy of 62.5 eV is needed to doubly ionize

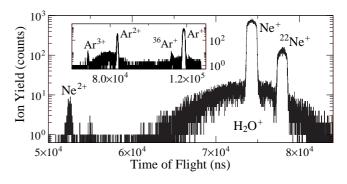


FIG. 1: Time-of-flight mass spectra of Ne⁽¹⁻²⁾⁺ and Ar⁽¹⁻³⁾⁺ ions (in the inset) at $I \approx 10^{12} \sim 10^{13}$ W/cm².

Ne atoms, which means that at least 2 photons have to be absorbed at the present photon energy of 38.8 eV. If, however, during the laser pulse the Ne atom gets first singly ionized by absorption of one photon and then, in a second step but within the same pulse, a second electron is removed from the relaxed Ne⁺ ground state, one more photon is needed requiring a total of 3 photons. SI can also proceed via two-photon absorption, if the first photoemission leaves the Ne⁺ in an excited shake-up state. From previous studies such shake-up satellites are known to contribute a few percent of the main line. According to lowest-order perturbation theory, expected to be valid at $I < 10^{15}$ W/cm² [16], the ionization yield should increase with I as $Y = \sigma_n \cdot I^n$, where σ_n is the generalized *n*-photon cross-section and *n* is the number of photons needed for ionization.

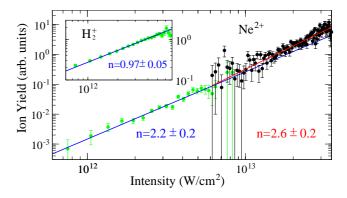


FIG. 2: Ion yield of Ne²⁺ and H₂⁺ (inset) as a function of *I*. Solid black and green points are the present data for two different sizes of the light collimation exit-slit. Solid blue and red lines are fits by $\log Y = n \cdot \log I + \log \sigma_n$ to two sets of data yielding $n = 2.2 \pm 0.2$ and $n = 2.6 \pm 0.2$. Single ionization of H₂ residual gas leading to the slope $n = 0.97 \pm 0.05$, provides further evidence for the linearity of our intensity scale.

The Ne²⁺ yield is analyzed as a function of I in Fig. 2. The blue and red curves, fitted to two sets of data, nicely describe the experimental intensity dependences of Ne²⁺ ions with the slopes $n = 2.2 \pm 0.2$ and $n = 2.6 \pm 0.2$, respectively. This indicates that double ionization of Ne is mainly induced via a 2-photon absorption at $I < 6 \times 10^{12}$ W/cm², clearly favoring

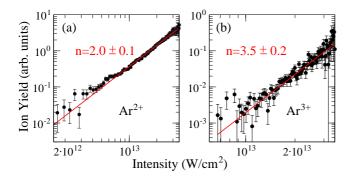


FIG. 3: Yield of Ar^{q+} ions as a function of I, q = 2 (a) and q = 3 (b); for details, see the caption of Fig. 2.

the NSI mechanism. The slope n = 2.6 found at higher intensities points to mixed NSI and SI transition contributions. As *I* increases, SI contributes more and more, which is expected theoretically for two-photon double ionization of He [17]. The ionization-rate ratio $P_{\text{Ne}^{2+}}/P_{\text{Ne}^{+}}$ agrees well with the measurements taken at the same photon energy by Sorokin *et al.* [18].

Similar conclusions can be drawn from the Ar^{3+} intensity dependence shown in Fig. 3(b). The slopes deliver the values $n = 2.0 \pm 0.1$ and $n = 3.5 \pm 0.2$ for Ar^{2+} and Ar^{3+} , respectively. According to the total ionization potentials for doubleand triple-ionization of Ar, NSI needs 2 or 3 photons and SI 2 or 4 photons for charge states q = 3 or 4, respectively. Analogous to the Ne data at high intensities, n = 3.5 indicates that both NSI and SI comparably contribute to the creation of Ar^{3+} . Since the same number of photons is needed to ionize Ar^{2+} for NSI as well as SI, the value $n = 2.0 \pm 0.1$, further confirms the reliability of the measured intensities and ionization rates. Our finding may not completely agree with recent results [10], where sequential ionization was observed to be the dominant mechanisms for MPI of Ar and Xe at comparable intensities but lower photon energies (12.7 eV).

In addition to the total ionization yield we were able to record the first recoil-ion momentum distribution for twophoton double ionization of Ne in Fig. 4. Shown are the ion-momentum components in the x-y-plane transverse to the light-propagation direction along z containing the polarization axis along y for various integration intervals along z as stated in the figure caption. In Fig. 4(a), the recoil-ion momentum distribution of Ne⁺ is observed to cluster on a circle with the radius $P_r = (P_x^2 + P_y^2)^{1/2} = 1.05$ a.u. as expected for absorption of one single photon. Ions produced by two-photon or second-harmonic photon absorption are clearly absent, expected to occur at the large circle at $P_r = 2.0$ a.u. in Fig. 4(a). This result, obtained with a statistical significance of about 10^{-2} relative to one-photon single ionization, agrees with the measured ion-yield dependence in Fig. 2.

The Ne²⁺ pattern, representing the vector sum-momentum distribution of the two electrons, is shown in Fig. 4(b) and (c) for $I < 3 \times 10^{12}$ W/cm² and $I > 2 \times 10^{13}$ W/cm², respectively. For NSI, the electron excess energy is ≈ 15 eV resulting

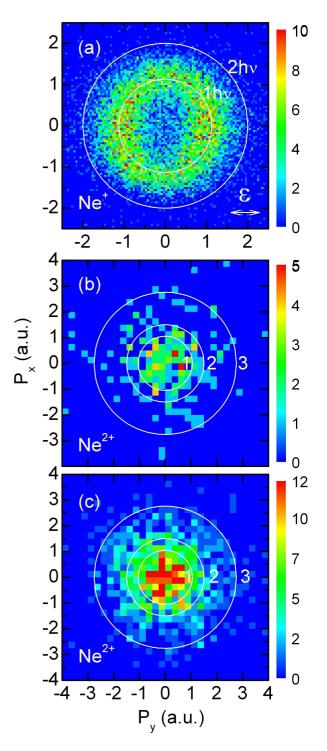


FIG. 4: Density plot of recoil-ion momentum distributions for Ne⁺ (a) and Ne²⁺ (b) and (c). The solid curves indicate the momentum positions for one and two photon absorption in (a) as well as for NSI and SI processes in (b) and (c) for various emission mechanisms, see text. Ne⁺ momenta are integrated over events with $|P_z| \leq 0.1$ a.u. at low intensities corresponding to the green points in Fig. 2. Ne²⁺ distributions integrate all events along the *z*-direction at $I < 3 \times 10^{12}$ W/cm² (b), and $I > 2 \times 10^{13}$ W/cm² (c). The polarization of the light is along the *y*-axis.

in $P_r \leq 1.5$ a.u. indicated as the circle 2 in Fig. 4 (b) and (c). For SI, i.e. absorption of three photons, the maximum twoelectron sum-momentum (emission into the same direction) can be 2.76 a.u. plotted as the circle 3. Denoting C_i to represent the numbers of events inside the circle i (i = 1, 2, 3), we might draw conclusions about the contribution of various processes. Thus, the rate $(C_3 - C_2)/C_3$ of events in Fig. 4(b) point to a 29% SI contribution and a $C_2/C_3 = 71\%$ contribution of NSI inside the circle 2. The observation of both NSI and SI in the ion-momentum spectra at $I < 3 \times 10^{12}$ W/cm² is in qualitative agreement with the intensity dependence presented in Fig. 2. The NSI is certainly the dominant transition process while SI is active simultaneously, though with a lower probability.

For higher light intensities, i.e., at $I > 2 \times 10^{13}$ W/cm², $(C_3 - C_2)/C_3$ increases to a 35% contribution, whereas C_2/C_3 slightly decreases to 65%, a clear indication that the SI-contribution increases with *I*, which supports the findings in the intensity dependence above. Note that due to present momentum resolution (≈ 0.8 a.u.) and overlapping emission patterns for NSI and SI, it is impossible to absolutely separate these two transition processes by measuring recoil-ion momenta alone. Thus, $C_3 - C_2$ could also include some events of NSI and vice versa.

The circle 1 in Figs. 4 (b) and (c) with a radius of 1.05 a.u. contains the events for two-electrons ejected most likely into opposite hemispheres assuming similar energies and amounts to $C_1/C_2 = 66\%$ of the NSI contribution in Fig. 4(b). For two-photon double ionization the angular momentum constraints for the relative emission of the two electrons are much more relaxed as compared to single-photon double ionization. The observed pattern shows a clear maximum at zero momentum corresponding to a kinematics where both electrons are emitted with similar energies into roughly opposite directions. For single photon double ionization this configuration is suppressed and even forbidden for equal energy sharing if the Ne^{2+} is in the electronic ground state (even symmetry). Single-photon double ionization of He, depicted in Fig. 1 of Ref. [19], shows a pronounced double-hump structure in the recoil-ion momentum distribution reflecting the dipole emission characteristics as a result of the angular momentum selection rules, in clear disagreement with the present findings. Back-to-back emission for NSI supports theoretical predictions for He [20], where this geometry was found to represent the main decay channel.

In summary, we studied few-photon multiple ionization of Ne and Ar using FLASH photons in the VUV-regime. The light-intensity dependence for ion-yields confirmed double ionization of Ne to be mainly due to two-photon nonsequential ionization at $I < 6 \times 10^{12}$ W/cm². As I increases, SI transitions for Ne²⁺ as well as Ar³⁺ production become more pronounced. Recoil-ion momentum distributions of Ne²⁺ confirm these observations and, for the first time, provide evidence for a sizeable contribution of events, where two ejected electrons widely share the excess energy available and, thus, are emitted with similar energies, into back-to-back directions. With the increase of the FLASH pulse repetition rate achieved recently, fully differential measurements are deeming at the horizon.

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- [1] O. Schwarzkopf et al., Phys. Rev. Lett. 70, 3008 (1993).
- [2] Th. Weber et al., Nature(London) 431, 437 (2004).
- [3] J. Briggs and V. Schmidt, J. Phys. B 33, R1 (2000).
- [4] A. Becker and F. H. M. Faisal, J. Phys. B 38, R1 (2005).
- [5] J. S. Parker et al., Phys. Rev. Lett. 96, 133001 (2006).
- [6] V. Ayvazyan et al., Eur. Phys. J. D. 37, 297 (2006).
- [7] J. Ullrich et al., Rep. Prog. Phys. 66, 1463 (2003).
- [8] T. Laarmann et al., Phys. Rev. A 72, 023409 (2005).
- [9] Y. Nabekawa et al., Phys. Rev. Lett. 94, 043001 (2005).
- [10] H. Wabnitz et al., Phys. Rev. Lett. 94, 023001 (2005).
- [11] R. Santra and C. Creene, Phys. Rev. A 70, 053401 (2004).
- [12] E. P. Benis et al., Phys. Rev. A 74, 051402(R) (2006).
- [13] D. M. P. Holland *et al.*, J. Phys. B **12**, 2465 (1979).
- [14] G. V. Marr and J. B. West, Atom. Data Nucl. Data Tables 18, 497 (1976).
- [15] S. Düsterer et al., Opt. Lett. 31, 1750 (2006).
- [16] M. Protopapas et al., Rep. Prog. Phys. 60, 389 (1997).
- [17] P. Lambropoulos et al., Phys. Rev. A 72, 013410 (2005).
- [18] A. Sorokim et al., Phys. Rev. Lett. (submitted) (2006).
- [19] A. Knapp et al., J. Phys. B 55, L521 (2002).
- [20] S. X. Hu et al., J. Phys. B 38, L35 (2005).