

**Transition from correlated to single-active-electron excitation in strontium nonlinear ionization**A. Dimitriou<sup>1,\*</sup>, V. Lorient<sup>2</sup>, A. Marciniak<sup>2</sup>, T. Barillot<sup>2</sup>, S. Danakas<sup>1</sup>, F. Lépine<sup>2</sup>, C. Bordas<sup>2</sup>, and S. Cohen<sup>1,†</sup><sup>1</sup>*Atomic and Molecular Physics Laboratory, Physics Department, University of Ioannina, 45110 Ioannina, Greece*<sup>2</sup>*Université de Lyon, CNRS, UMR5306, Institut Lumière Matière, 69622 Villeurbanne, France*

(Received 30 October 2021; accepted 25 April 2022; published 9 May 2022)

The nonlinear single and double ionization of Sr atoms as a result of their interaction with 800-nm Ti-sapphire laser pulses of 25-fs duration is experimentally investigated within the laser intensity range  $I = 3 \text{ TW cm}^{-2}$ – $30 \text{ TW cm}^{-2}$ . The latter range includes the over-the-barrier intensities for both Sr and  $\text{Sr}^+$ , while the corresponding Keldysh adiabaticity parameters are larger than unity. Nevertheless, for  $I \geq 24 \text{ TW cm}^{-2}$  the recorded photoelectron-energy spectra show evidence of the (partial) applicability of tunneling or over-the-barrier ionization concepts, whereas below this value they can be discussed in terms of multiphoton ionization processes. More importantly, the data additionally point towards a further division of the multiphoton regime into two subranges separated roughly at  $I \sim 4 \text{ TW cm}^{-2}$ . Above this value above-threshold-ionization structures and the single-active-electron picture dominate. On the contrary, at the lowest laser intensities the photoelectron spectra and angular distributions reveal single ionization pathways where the remaining ion is left into excited states. These pathways also require photon absorption in the continuum, which in the case of strontium and other alkaline-earth-metal atoms is structured by the presence of doubly excited states which are embedded in it. Hence, in this case ionization proceeds via quasiresonant ionization ladders formed by these doubly excited states which are dominated by the interaction between the two valence electrons of Sr. We discuss in detail the above rich phenomenology and relevant ionization mechanisms and propose possible directions of further work towards the elucidation of the role of configuration interaction in ionization processes.

DOI: [10.1103/PhysRevA.105.053106](https://doi.org/10.1103/PhysRevA.105.053106)**I. INTRODUCTION**

The nonlinear (single, double, or even multiple) ionization of multielectron atoms is a subject of high interest in photo-physics as it questions the nature of electron correlation. Early investigations in intense laser-atom interaction concerned alkaline-earth-metal atoms [1–12] ionized by long laser pulses ( $\sim$ ns or  $\sim$ ps) at moderate intensity ( $\leq 10 \text{ TW cm}^{-2}$ ). They were soon followed by experiments on noble gases [13], the latter offering much higher ionization limits that are capable of withstanding higher laser intensities available with ultra-short ( $\sim$ fs) laser pulses. With the emergence of strong-field atomic physics, nonlinear ionization was categorized with the help of simple concepts such as the Keldysh adiabaticity parameter [14],  $\gamma = [V_I/(2U_p)]^{1/2}$ , expressed via the target's ionization potential  $V_I$  and the ponderomotive energy  $U_p = I/4\omega^2$  (in atomic units) of the laser field of frequency  $\omega$  and intensity  $I$  [15]. According to such a categorization, the condition  $\gamma > 1$  characterizes the perturbative multiphoton ionization (MPI) regime leading to above-threshold ionization (ATI) [16]. In ATI the ionized electron absorbs photons in excess of the minimum number required to overcome the ionization threshold. On the other hand the condition  $\gamma < 1$  is

fulfilled at higher laser intensity and characterizes the non-perturbative tunnel ionization regime [15]. Further increase of the intensity leads to over-the-barrier ionization (OBI) [15], where above a critical intensity  $I_{\text{OBI}}$  the light field completely suppresses the Coulomb barrier. In fact, this simple description is often insufficient because the nonhomogeneous spatiotemporal distribution of the laser intensity may mix the different regimes [17]. It is to be further emphasized that the above description relies essentially on a single-active-electron picture [18]. Within this latter context, alkali metal atoms having a single-valence electron outside closed subshells were recently used as model target systems [19–27]. For these low- $V_I$  atoms  $I_{\text{OBI}}$  is sufficiently low and leads to  $\gamma(I_{\text{OBI}}) > 1$ . Surprisingly, under such an apparently conflicting condition the ionization of these systems could be perfectly described in terms of MPI [20,25].

Alkaline-earth-metal atoms are also low- $V_I$  systems having two-valence electrons outside a closed-subshell ionic core (ground-state configurations  $[\dots]n_g s^2$ , where  $[\dots]$  denotes the core). Thus, they appear to be suitable for studying the role of multielectron effects which are at play in nonlinear ionization. Indeed, earlier long-pulse and low-intensity investigations [5,7,9–12] already pointed out that the single-active-electron assumption is frequently invalid for these systems. Excitation of the two-valence electrons results in dense manifolds of doubly excited states located near their first  $[\dots]n_g s$  ionization thresholds and all the way up to their double-ionization limits. These states were found to act as resonant or near-resonant

\*Present address: Institute of Nanoscience and Nanotechnology, NCSR “Demokritos,” Aghia Paraskevi, Athens, Greece.

†scohen@uoi.gr

intermediate levels and form excitation ladders that led to the absorption of a number of photons within the (structured) continuum. This process differs qualitatively from the single-active-electron ATI process mentioned earlier. In addition, ionization via such ladder schemes was found to create populations of *excited* states of the singly charged ions [5]. This effect constitutes a typical signature of the interaction between the two-valence electrons and the consequent extensive configuration interaction (CI) and mixing between (quasidiscrete and continuum) doubly excited configurations [28]. Despite this fact, “direct” two-electron ejection was never evidenced in alkaline earths under these conditions. Nevertheless, the production of excited ionic-level populations facilitated their sequential double ionization where atoms are ionized first and the resulting singly charged ions are further ionized within the same laser pulse [1–12].

Until recently nonlinear ionization of alkaline earths using ultrashort and intense fs pulses was rather rarely investigated (either theoretically [29–31] or experimentally [32–36]) with a particular focus on the Mg atom. Indeed, the first ionization threshold of Mg ( $\approx 7.6$  eV) was anticipated to be sufficiently high for allowing a meaningful comparison with the results of short-pulse nonlinear ionization experiments in noble gases [33]. The measured ATI electron spectra in Mg could be interpreted in terms of single-active-electron ionization, while a weaker ATI series originating from CI-driven ionization and leaving the ion to its excited  $3p$  state was unfortunately blended with the sequential double-ionization signal [32]. Further investigations concerned the resonant structures exhibited by the ATI peaks [35] and the nonsequential double ionization [34]. More recently, ionization induced by recollision processes was investigated at longer wavelength (1–2  $\mu\text{m}$ ) [37–39] in Mg and heavier alkaline-earth-metal atoms. The general conclusion drawn from these studies is that nonsequential double ionization proceeds via the excited ionic states which are pumped by the impact of the returning electron, but the atom-specific doubly excited spectrum does not seem to play any role. Consequently, the implication of these CI-mediated states to the overall laser-induced dynamics throughout from MPI to OBI still remains an open issue requiring further study.

The purpose of the present work is to shed more light on the above issue, by investigating the nonlinear single and double ionization of Sr using intense  $\approx 800$ -nm and  $\approx 25$ -fs laser pulses and a velocity-map imaging spectrometer (VMIS). Strontium atom is chosen because earlier long-pulse studies showed that CI-mediated above-threshold ionization is much stronger for it [10–12] as compared to other alkaline-earth-metal atoms [7,9] under similar excitation conditions. The recorded photoelectron energy spectra (PES) and photoelectron angular distributions (PADs) show indeed such evidence. However, as the laser intensity is increased we first observe a transition towards single-active-electron ATI structures and as it is increased even more we observe another transition towards the tunneling regime onset. Although this is a by-product of our principal investigation, the latter MPI-to-tunneling ionization transition is also discussed here in connection with the aforementioned recent studies in alkali atoms under conflicting conditions. These results offer unexplored perspectives for studying electron correlation in

the ultrafast regime and may prove useful for putting into stringent test future relevant theoretical models.

## II. EXPERIMENT SETUP AND PROCEDURE

The experimental setup consists of two differentially pumped vacuum chambers connected via an electronically controlled valve. The first chamber encloses an electrically heated stainless-steel oven filled with strontium pellets and heated to  $\approx 550$  °C. The produced Sr vapor passes through a 0.5-mm-diameter hole and forms a thermal atomic beam entering the second chamber when the valve is open. This laser-atom interaction chamber is pumped by a 250-l/s turbomolecular pump–rotary pump system maintaining a background pressure of  $\sim 10^{-7}$  mbar.

Strontium atoms interact with laser radiation delivered by a Ti: sapphire laser (Coherent, Legend Elite Duo USX) producing pulses of  $\approx 800$ -nm central wavelength,  $\approx 52$ -nm bandwidth, and  $\approx 25$ -fs pulse duration, while operating at 5-kHz repetition rate. The linearly polarized laser beam emerging from this system passes through an assembly composed of an achromatic half wave plate and a (dielectric thin-film multilayer) linear polarizer for setting its polarization parallel to the electron detector plane (see below) and for intensity control. Subsequently the beam is focused into the interaction region by a 25-cm focal length lens.

The atomic and laser beams are perpendicular to each other, as well as to the axis of the VMIS. The latter is based on a standard three-electrode design [40], where photoionization takes place in the center between a solid repeller plate and an extractor plate 20 mm apart and with a 20-mm central hole. These electrodes are followed by an equally spaced grounded third electrode identical to the extractor plate. Typically, electrode voltages are  $V_{\text{rep}} \approx -3.5$  kV and  $V_{\text{ext}} \approx -2.5$  kV, in order to ensure proper VMIS focusing conditions [40,41]. The electric field created by the three electrodes accelerates photoelectrons towards the end of a 15-cm-long field-free drift tube where they are detected by a two-dimensional position-sensitive detector. The latter is made of a tandem Hamamatsu F2225-21P microchannel plate (MCP) assembly and a P42 phosphor screen. The entire spectrometer is shielded against parasitic magnetic fields by means of a double  $\mu$ -metal layer. A charge-coupled device camera (Allied Vision, Marlin F146B) records the 2D distribution of light spots on the screen. Recorded images are transferred to a PC, where they are accumulated over several hundred thousand laser shots.

The processing of raw images is performed off-line. The PES and PADs are obtained from the recorded 2D projections. For cross-checking purposes we employed two inversion procedures based on Vrakking’s method [42] and the so-called “polar onion peeling” one [43]. Both methods produced identical results within experimental resolution and the PADs presented in this work are those obtained by Vrakking’s method. Electron-energy calibration is achieved via the observed ATI electron peaks. Judging from the measured smallest energy difference between closely spaced electron peaks and their widths, the resolution of the spectrometer is better than  $\approx 0.1$  eV for electron energies on the order of 1 eV. As for the detection of produced ions, it is achieved by

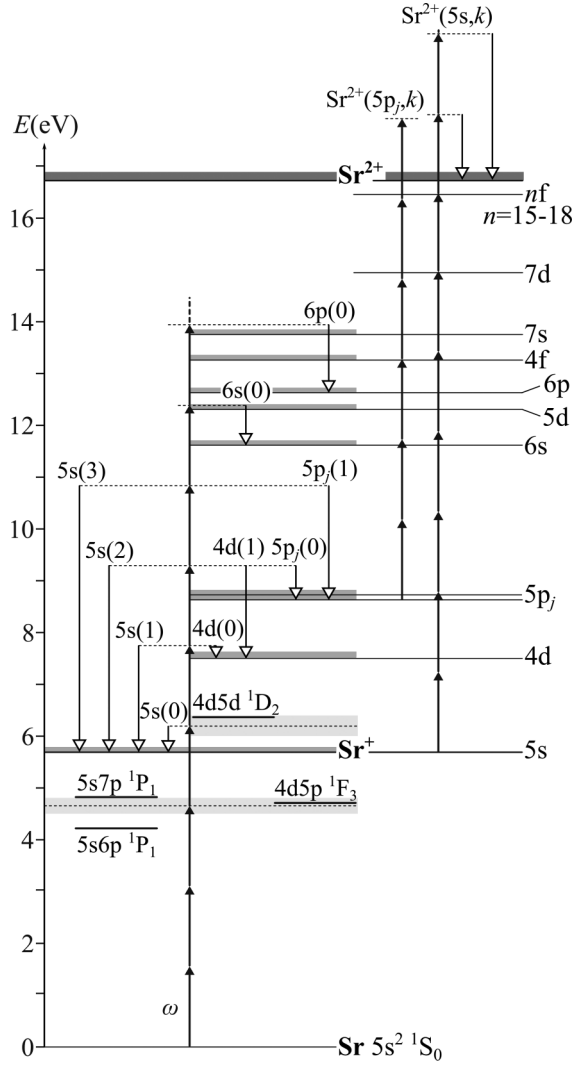


FIG. 1. Partial, in-scale, energy-level diagram of Sr, depicting the most relevant atomic and ionic levels as well as the possible multiphoton single and double-ionization pathways. Dark gray shadings denote the ionization threshold and continua associated with each ionic level and the double-ionization limit. Light gray shading denotes the three- and four-photon effective laser bandwidth [49]. For the labeling of ionization processes see the text [and Eqs. (3)–(9)].

simply reversing the polarities of the voltages applied to the interaction-region electrodes of the VMIS and operating the instrument as a simple ion time-of-flight mass spectrometer. Finally, the laser-beam intensity calibration is accomplished by employing this ion detection operating mode and measuring the ratio between the single and double-ionization yield of Xe [44] in a separate study. We estimate that our reported laser intensities are accurate to within 10%.

### III. EXCITATION AND IONIZATION PATHWAYS

Before presenting our experimental data it will be helpful to discuss first the relevant excitation and ionization pathways. These are schematically shown in the scaled (though partial) energy-level diagram of Fig. 1 without considering any ponderomotive and/or dynamic (AC) Stark level shifts.

We initially restrict ourselves to the processes where the atom, initially in its  $5s^2 \ ^1S_0$  ground state, absorbs a number of photons and ionizes, leaving the singly charged ion to a given  $n_i \ell_i$  level. By assuming strictly nonresonant ionization conditions and that the atomic  $5s^2$  ground state is AC Stark shifted by  $\Delta E_{5s^2}$  while the  $n_i \ell_i$  limit  $V_{1, n_i \ell_i}$  is shifted by  $U_p$  [45], the kinetic energy of the electron released by such a process may be approximated by

$$\varepsilon_{n_i \ell_i, N} = N\omega - V_{1, n_i \ell_i} - (U_p - \Delta E_{5s^2}). \quad (1)$$

Just like the ponderomotive energy [given by  $U_p(\text{eV}) = 5.95 \times 10^{-2} I$  (TW cm $^{-2}$ ) for the wavelength employed here], the quantity  $\Delta E_{5s^2}$  is expected to vary linearly with laser intensity  $I$ . As for the integer  $N$ , it stands for the total number of absorbed photons out of the ground state,

$$N = N_{n_i \ell_i} + k, \quad k = 0, 1, 2, \dots, \quad (2)$$

where  $N_{n_i \ell_i}$  is the minimum number required to ionize the atom with respect to the  $n_i \ell_i$  threshold and integer  $k$  describes the ATI process associated with this threshold. To facilitate our discussion we label the above-described process as  $n_i \ell_i(k)$ , and the pathways that are or could be of relevance to the present work are summarized in the following equations (3)–(9), most of them drawn also in Fig. 1:

$$5s(k): \quad \text{Sr}(5s^2) + (4+k)\omega \rightarrow \text{Sr}^+(5s) + e(0.51 \text{ eV} + k\omega), \quad (3)$$

$$4d(k): \quad \text{Sr}(5s^2) + (5+k)\omega \rightarrow \text{Sr}^+(4d) + e(0.23 \text{ eV} + k\omega), \quad (4)$$

$$5p_j(k): \quad \text{Sr}(5s^2) + (6+k)\omega \rightarrow \text{Sr}^+(5p_j) + e[0.57(j=3/2)/0.67(j=1/2) \text{ eV} + k\omega], \quad (5)$$

$$6s(k): \quad \text{Sr}(5s^2) + (8+k)\omega \rightarrow \text{Sr}^+(6s) + e(0.79 \text{ eV} + k\omega), \quad (6)$$

$$6p(k): \quad \text{Sr}(5s^2) + (9+k)\omega \rightarrow \text{Sr}^+(6p) + e(1.32 \text{ eV} + k\omega), \quad (7)$$

$$4f(k): \quad \text{Sr}(5s^2) + (9+k)\omega \rightarrow \text{Sr}^+(4f) + e(0.69 \text{ eV} + k\omega), \quad (8)$$

$$7s(k): \quad \text{Sr}(5s^2) + (9+k)\omega \rightarrow \text{Sr}^+(7s) + e(0.20 \text{ eV} + k\omega), \quad (9)$$

where  $\omega \approx 1.55$  eV. The process corresponding to Eq. (5) is denoted as  $5p_j(k)$  and the  $\approx 0.1$ -eV spin-orbit splitting is taken into account because it is comparable to the energy resolution of our spectrometer. Note further that the  $k=0$  electron energies reported in the above equations were computed without considering any kind of shift. If  $U_p$  and  $\Delta E_{5s^2}$  are considered in Eq. (1), the above given electron energies decrease linearly with  $I$  and Fig. 1 turns out to be somewhat misleading. For example, by ignoring  $\Delta E_{5s^2}$  to a first approximation, Eq. (1) predicts that for the lowest intensity employed here

the  $5d(0)$  channel [ $\text{Sr}(5s^2) + 8\omega \rightarrow \text{Sr}^+(5d) + e$ ] is expected to be *closed*, that is,  $\varepsilon_{5d,8}(I) \leq 0$  [45]. For this reason we do not consider this process in the above equations. In addition, for the same intensity the  $4d(0)$  and  $7s(0)$  channels are about to be closed. It should be stressed, however, that high-lying  $[...n_i\ell_i n\ell]$  series of Rydberg states and the  $n_i\ell_i$  thresholds these series converge to are expected to suffer (approximately) the same  $U_p$  shift. This circumstance has two consequences. First, even above the channel closing intensity these excited levels may sustain ionization by gradually upshifting to resonance within the time development of the laser pulse (the so-called Freeman resonances [46]). Second, either below or above the closing intensity the energies of electron peaks corresponding to Freeman resonances are approximately independent of  $I$ . Then, Eq. (1) is invalid and the initial level AC Stark shift and  $U_p$  may solely affect the threshold intensity where the resonance condition is satisfied.

Leaving aside the missing shifts, the atomic ionization pathways of Fig. 1 are drawn in a way that allows their broad classification into typical ATI processes and nontypical absorption in the structured continuum. Consider, for example, the  $5s(k > 0)$  pathway belonging to the former category, where one optically active electron absorbs photons above the first ionization threshold. Then, ionization occurs with a certain probability at each absorption step, while the singly charged ion is always left unexcited, that is, the  $5s$  ground-state electron is a spectator. In other words, this ATI channel involves only  $5s\varepsilon\ell$  atomic continuum configurations, with  $\varepsilon$  and  $\ell$  determined by energy conservation and angular momentum dipole selection rules, respectively (recall that absorption of linearly polarized radiation obeys the  $\Delta m_J = 0$ ,  $\Delta J = \pm 1$  and parity change dipole selection rules per each absorbed photon).

Consider next those processes where atomic MPI results in an excited singly charged ion, as described by Eqs. (4)–(9). This mechanism strongly depends on the existence of energy-level structure in the energy ranges reached after absorption of  $m$  photons from the atomic ground state ( $m = 1, 2, 3$ , etc.) and, therefore, its understanding requires the examination of this structure first. Let us begin by noting that there are no resonant or near-resonant states at the first and second photon level [47,48]. On the contrary, by taking into account the *effective* three-photon laser bandwidth [49] we find the  $4d5p\ ^1F_3$  and  $5s7p\ ^1P_1$  three-photon near-resonant states, both lying slightly *above* the central laser frequency (see Fig. 1). In fact the  $\ ^1F_3$  level lies below the full  $5snf\ ^1F_3$   $n \geq 4$  Rydberg series and has both  $4d5p$  ( $\sim 25$ – $30\%$ ) and  $5snf$  character [50]. Similarly, the  $5s7p\ ^1P_1$  state also comprises about 13% of doubly excited  $4d5p$  character [48,50]. Passing now to the four-photon level, we find that the available  $5s\varepsilon\ell$  continua with  $\ell = J = 0, 2$ , and  $4$  are mixed by CI with a number of near-resonant autoionizing states. Specifically, the effective laser bandwidth comprises the whole  $4d5d$  doubly excited configuration [48] and notably the  $\ ^1D_2$  level shown in Fig. 1. The  $J = 0, 2$  states are additionally expected to have an important admixture of  $5p^2$  character [51]. As for the range above the  $4d$  threshold (vicinity of the fifth photon), it is dominated by the spectrally broad low-lying members of  $[5pns]_{J=1}$  [52] and  $[5pnd]_{J=1,3}$  [53] series. Lastly, no doubly excited states are expected in the vicinity of

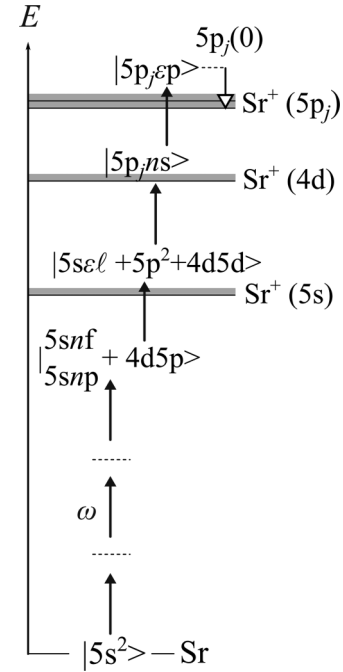


FIG. 2. Schematic diagram of a ladder excitation scheme describing a possible (out of several)  $5p_j(0)$  ionization pathway producing excited ions [Eq. (5)]. Gray shading denotes the ionization thresholds and continua associated with  $5s$ ,  $4d$ , and  $5p_j$  ionic levels. At this particular scheme each ladder step involves single-electron transitions but, overall, both valence electrons are excited. Such ladders necessitate the existence of electron-electron correlation leading to strong configuration interaction and level structure embedded in the continuum, the latter characterizing Sr and the other alkaline-earth-metal atoms. Note that the experiment probes the population created in the excited ionic state but not the exact ladder scheme.

the sixth photon, while the seventh and higher photon ranges are largely unexplored and the data are fragmentary [54].

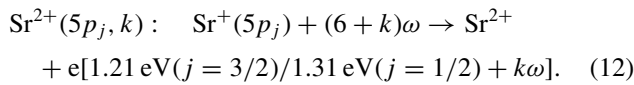
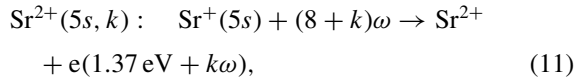
Having the above presentation in mind one may anticipate that the aforementioned configuration mixings may trigger excitation ladders connecting doubly excited states at each step. As an example, consider the process

$$\begin{aligned} |5s^2\rangle &\xrightarrow{+3\omega} |\dots + 4d5p\rangle \xrightarrow{+\omega} |\dots + 4d5d + 5p^2\rangle \\ &\xrightarrow{+\omega} |5p_jns\rangle \xrightarrow{+\omega} |5p_j\varepsilon p\rangle, \end{aligned} \quad (10)$$

which is also depicted schematically in Fig. 2 and which describes a possible  $5p_j(0)$  ionization pathway producing  $\text{Sr}^+$  excited to its  $5p_j$  state. Although in this example each ladder step was chosen to mainly involve single-electron transitions, overall both valence electrons are excited. Therefore, ladders of this type cannot be formed in the absence of electron-electron correlation and level structure embedded in the continuum, this situation being a specific feature of alkaline-earth elements. As a reminiscence of these facts the relevant ionization pathways are hereafter labeled as CI-ATI. Nevertheless, typical ATI is also possible here, leaving the ion always in the same excited state [4]. Thus, the  $5p_j(1)$  pathway

in Fig. 1 involves both CI-ATI and ATI. In general, these two processes may run in parallel, each delivering a quite specific PES that constitutes its fingerprint.

Finally, in order to discuss double ionization we need to emphasize that the reference to electron correlation notions made here differs qualitatively from the one invoked for the description of nonsequential double-electron ejection under the influence of strong, short-pulse and long-wavelength laser fields [37–39]. In the latter case “correlation” refers to collisional effects between two “quasifree” electrons that are induced by the interacting laser field. By contrast, the presently discussed CI-ATI relies on the atom-specific bound and autoionizing energy-level structure, although this structure can, up to a certain extent, be perturbed by the laser field. Moreover, while CI-ATI may also lead to nonsequential (or direct [5]) double ionization, all the processes described by Eqs. (3)–(9) lead to sequential multiphoton double ionization. We denote double-ion production out of a given initial ionic level as  $\text{Sr}^{2+}(n_i \ell_i, k)$  and Fig. 1 shows the examples of the following  $\text{Sr}^{2+}(5s, k)$  and  $\text{Sr}^{2+}(5p_j, k)$  pathways:



Like Eqs. (3)–(9), we include here the possibility of ATI, while the reported  $k=0$  electron energies apply in the absence of ponderomotive or AC Stark shifts and Freeman resonances.

## IV. RESULTS AND DISCUSSION

### A. $\text{Sr}^+$ and $\text{Sr}^{2+}$ yields

The  $\text{Sr}^+$  and  $\text{Sr}^{2+}$  ion yields as a function of laser intensity are presented in Fig. 3. We note first that below saturation the singly charged ion yield follows an  $I^{K_+}$  power law with an apparent slope of  $K_+ \approx 4$ . Saturation occurs at  $\sim 6 \text{ TW cm}^{-2}$ , i.e., at a value which is higher than the one reported earlier for ionization studies via long ( $\sim \text{ns}$ ) pulses of similar wavelength and processes of the same minimum nonlinearity of ionization [10]. This is to be expected since in the present work pulses of shorter duration (fs) are employed. As for the  $\text{Sr}^{2+}$  yield, it does not saturate up to the highest intensity of the graph; it exhibits an ionization order of nonlinearity of  $K_{2+} \approx 3$  and it appears to be detectable at intensities equal to or greater than the above  $\text{Sr}^+$  saturation intensity. This is a signature of a sequential double-ionization mechanism. It additionally suggests that for  $I < \sim 5 \text{ TW cm}^{-2}$  we can ignore, to a first approximation, double-ionization processes for the analysis of the PES.

Although not related to our main focus on the study of CI-ATI processes, it would be interesting at this point to open a parenthesis and briefly discuss the connection of the presently recorded data with the aforementioned alkali atom ionization studies. In order to implicate over-the-barrier concepts to this discussion, let us first use the most frequently employed 1D expression for the critical intensity  $I_{\text{OBI}}$  which

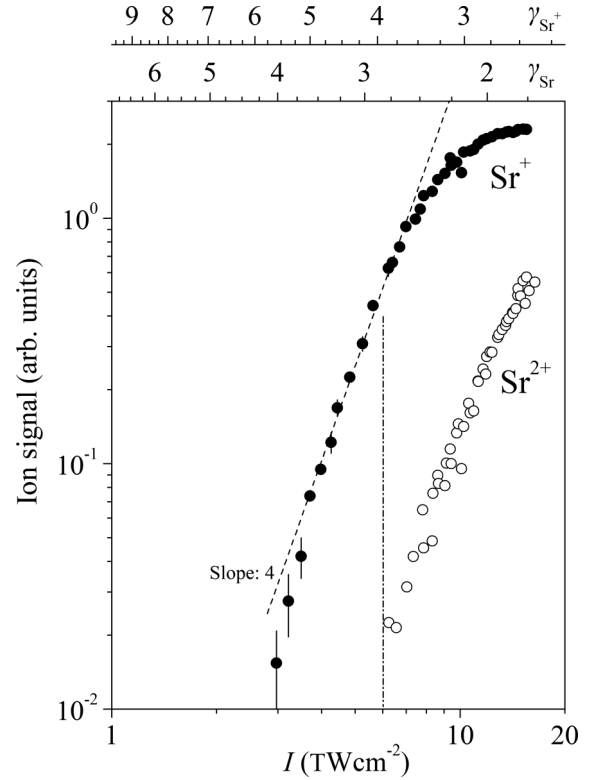


FIG. 3. Log-log plots of the measured singly and doubly charged Sr yields as a function of laser intensity (lower  $x$  axis) and as a function of the atomic and ionic Keldysh parameters  $\gamma_{\text{Sr}}$  and  $\gamma_{\text{Sr}^+}$ , respectively (upper  $x$  axes). Whenever the  $y$  axis uncertainties are not indicated they are on the order of the symbol size. A dashed straight line corresponding to an  $I^4$  intensity dependence is matched to the unsaturated part of the  $\text{Sr}^+$  curve. The latter is separated from the saturated one by the vertical dashed-dotted line. A quantifiable  $\text{Sr}^{2+}$  signal appears for intensities above the  $\text{Sr}^+$  saturation intensity. A fit of the doubly charged ion signal as a function of  $I$  yields an  $I^{3.0 \pm 0.1}$  intensity dependence (not shown).

writes (in atomic units) [15,20] as

$$I_{\text{OBI}} = \left[ \frac{\kappa^4}{16Z_{\text{eff}}} \right]^2, \quad (13)$$

where  $Z_{\text{eff}}$  is the effective charge of the residual ion and  $\kappa = \sqrt{2V_1}$  (atomic units). It is presently accepted that this expression underestimates  $I_{\text{OBI}}$  [15]. Indeed, for Sr atom ( $V_1 = 5.7 \text{ eV}$ ,  $Z_{\text{eff}} = 1$ ) Eq. (13) predicts  $I_{\text{OBI}}(\text{Sr}) \approx 4 \text{ TW cm}^{-2}$ , which is close to the values reported for several alkali atoms with comparable  $V_1$ 's [19,22,25]. However, it lies below the saturation intensity of the  $\text{Sr}^+$  yield, contrary to the expectation that for  $I \geq I_{\text{OBI}}$  the ionization probability should be unity (saturated ionization process) [25]. Similarly, for  $\text{Sr}^+$  ( $V_1 = 11.03 \text{ eV}$ ,  $Z_{\text{eff}} = 2$ ) Eq. (13) predicts  $I_{\text{OBI}}(\text{Sr}^+) \approx 15 \text{ TW cm}^{-2}$ , again far below saturation. A presumably more accurate estimate for  $I_{\text{OBI}}$  may be obtained by adopting the less familiar 3D formula based on the parabolic coordinates  $(\xi, \eta, \varphi)$  [20,55] (in atomic units):

$$I_{\text{OBI},\eta} = \left[ \frac{\kappa^4}{16Z_{\text{eff}} - 8\kappa} \right]^2, \quad (14)$$

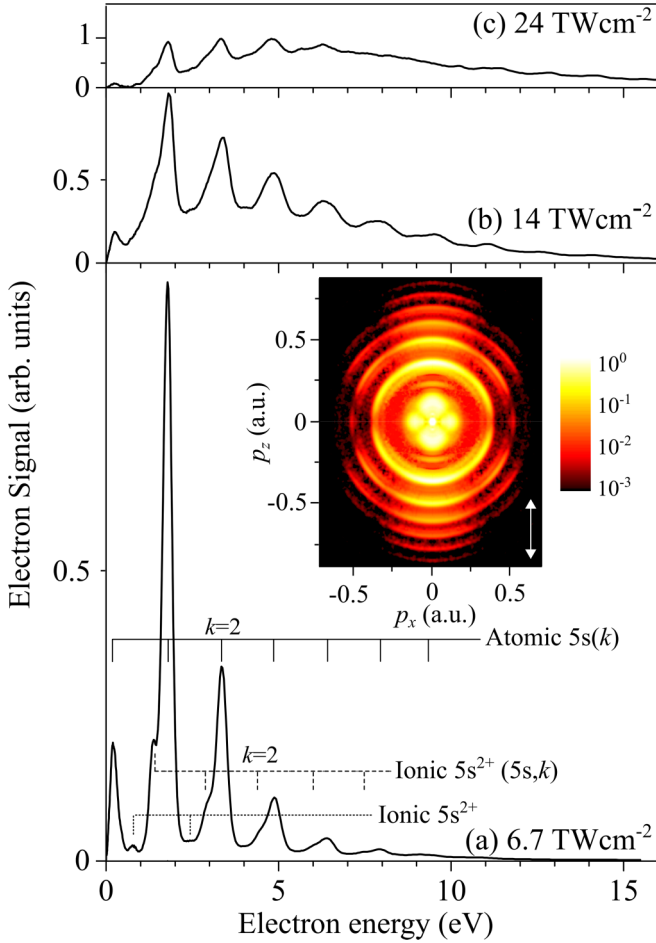


FIG. 4. Electron-energy spectra acquired at the laser intensity values specified at each plot. The maximum signal of each PES is set equal to unity. In (a) the members of the atomic and ionic ATI series discussed in the text are indicated. The corresponding inverted (and symmetrized) image given in the inset shows the photoelectron momentum distributions (in atomic units, a.u.)/PADs. For better visibility of its weak features the image is drawn in logarithmic color scale covering three orders of magnitude (see the given color bar). The white double-headed arrow shows the direction of the linear laser polarization. Note that in (b) the number of ATI members increases further and a background starts to build up, while this background dominates in (c) and the ATI structure is greatly washed out.

where the symbol  $I_{\text{OBI},\eta}$  is used for making the distinction from Eq. (13). Yet, both estimates rely on the static field concept. Using Eq. (14) we find  $I_{\text{OBI},\eta}(\text{Sr}) \approx 9 \text{ TW cm}^{-2}$  and  $I_{\text{OBI},\eta}(\text{Sr}^+) \approx 25 \text{ TW cm}^{-2}$ , which appear more compatible with the yields of Fig. 3 despite the  $\gamma > 1$  condition holding over the whole intensity range of the present experiment (see upper  $x$  axes of Fig. 3) and suggesting an MPI picture. A finer characterization of the ionization regime is to be provided by the following analysis of PES.

### B. Evolution of PES with laser intensity

Typical PES recorded within the  $4 \leq \gamma_{\text{Sr}} \leq 1.2$  and  $5.5 \leq \gamma_{\text{Sr}^+} \leq 1.7$  intervals are given in Figs. 4 and 5. Figure 4

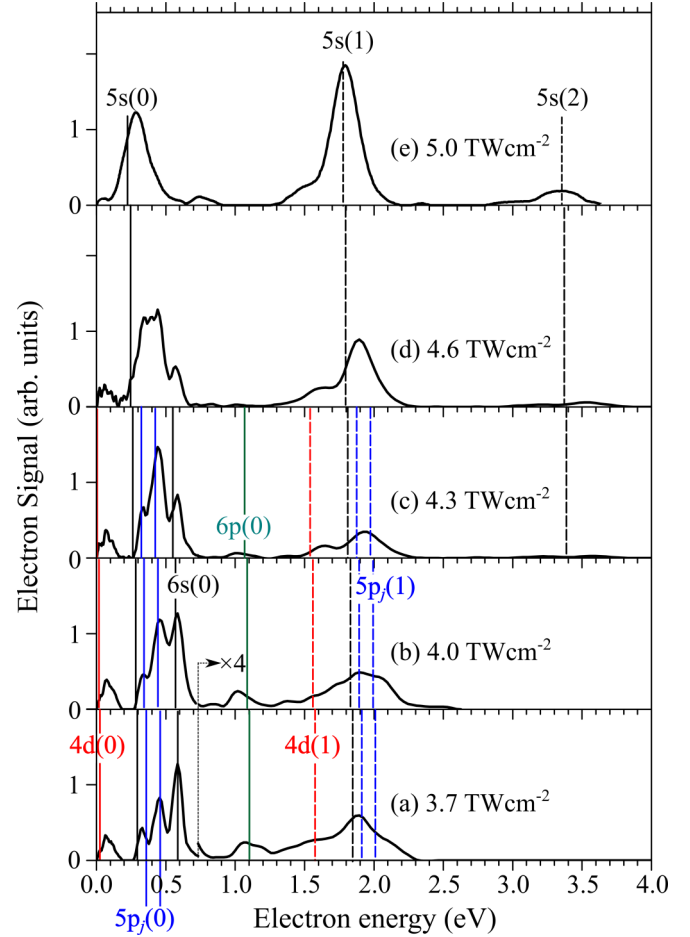


FIG. 5. Electron-energy spectra acquired at the lower intensity range 3.7–5  $\text{TW cm}^{-2}$ . The location of each spectrum along the  $y$  direction is proportional to the intensity of the particular recording which is specified at each plot. For facilitating the discussion given in the text the graph also includes the predictions of Eq. (1) for the single-ionization pathways of Eqs. (3)–(7), ( $k = 0$  full vertical lines,  $k > 0$  dashed vertical lines) after including only the ponderomotive shift  $U_p$ . The  $5s^2$  ground state AC Stark shift is ignored (i.e.,  $\Delta E_{5s^2} = 0$ ) because its inclusion using the existing theoretical value [57] considerably worsens the agreement with all experimental data. Furthermore, since within this intensity range double ionization is not apparent in Fig. (3), the relevant ionization processes are not included in the graph (for a more detailed discussion of this point see text).

presents a collection of the data corresponding to the higher intensity interval (roughly 7–24  $\text{TW cm}^{-2}$ ) where the singly charged  $\text{Sr}^+$  ion is saturated and double ionization is efficient ( $\gamma_{\text{Sr}} < 2.8$  and  $\gamma_{\text{Sr}^+} < 3.9$ ). Within this interval the PES evolve from typical single-active-electron ATI structures towards a mixed ATI-tunneling regime. In the following presentation the ATI series are discussed first because they will serve as a “reference” when seeking points of departure from their apparently simple structure. The points of departure from this reference at higher intensities concern the above-mentioned onset of tunneling which is discussed afterwards. Although a by-product of the present study with no connection to CI-ATI signatures, this mixed regime is here commented

upon for completeness and for making again a connection with earlier recent research devoted to strong field nonlinear ionization of the alkali atoms. As for the finally discussed points of departure from the typical ATI structures that are related to CI-ATI, i.e., to the main subject of interest in the present work, they are evident in the PES given in Fig. 5. This graph shows typical spectra concerning the lower-intensity interval, 3.7–5 TW cm<sup>-2</sup>, where the Sr<sup>+</sup> yield is unsaturated and Sr<sup>2+</sup> appears to be undetectable. Below, we analyze the spectra of these two graphs in more detail and invoke the relevant PADs when necessary, for shedding light on the involved ionization mechanisms.

### 1. Intensity range above the Sr<sup>+</sup> saturation intensity: ATI structures

We focus first on the reference curves, Figs. 4(a) and 4(b), whose main feature is the presence of several strong ATI peaks. The curve of Fig. 4(a) is recorded at  $I \approx 6.7$  TW cm<sup>-2</sup>, where the Sr<sup>2+</sup> signal is apparent. Therefore, it is reasonable to assume that the observed ATI series stem from both single and double-ionization processes (the latter contributing a much weaker signal to the total one). A careful look at the lower-energy and strongest ATI members reveals a double-peak structure that implies the simultaneous presence of two ATI series. Actually, two members of a third quite weak ATI series are also visible in the spectrum of Fig. 4(a) and the corresponding inverted VMI image given in its inset. Note further that within the examined intensity range the peaks of all three series remain practically unshifted with varying intensity. This peak stabilization cannot stem from a cancellation effect between  $U_p$  and initial state AC Stark shifts. For the atomic system this is ruled out because the low-energy ATI peaks do shift at lower intensities. By using theoretical values of dynamic dipole polarizabilities and corresponding AC Stark shifts [56] we verified that the ionic system suffers no such type of cancellation as well. In another scenario, since the Sr<sup>+</sup> yield is saturated, ionization could occur at the same intensity which is reached at the leading edge of the pulse and irrespective of its maximum value [19,20,25]. Nevertheless, this scenario is improbable for the spectra of Figs. 4(a) and 4(b) because within this intensity range the ATI structure continues to develop and the number of ATI peaks increases with  $I$ . Furthermore, this scenario is irrelevant for the ionic system (unsaturated Sr<sup>2+</sup> signal). Consequently, the effect has to be attributed to the presence of Freeman resonances whose energy location is indeed approximately independent of  $I$ .

Dealing with the atomic system first, we find that there is only one singlet three-photon Freeman resonance due to the bound  $5s6p\ ^1P_1$  state (see Fig. 1) that is compatible with the spectrum of Fig. 4(a) at the given laser intensity. The latter is very close to the  $I_{\text{thr}} \approx 7.2$ -TW cm<sup>-2</sup> threshold intensity of appearance for this resonance, as estimated by setting  $\Delta E_{5s^2} = 0$  and using the equation  $[E_{5s6p\ ^1P_1} + U_p(I_{\text{thr}})] = 3\omega$ . In fact, since experimentally this peak stabilizes at  $I \approx 6$  TW cm<sup>-2</sup>, it is reasonable to assume that the actual threshold is somewhat lower than the above value (as implied by the condition  $\Delta E_{5s^2} < 0$ —see the calculation of Ref. [57]). Furthermore, this resonance is expected to produce  $\approx 0.08$ -eV electrons, i.e., approximately equal to the energy of  $\sim 0.1$  eV correspond-

ing to the *nonresonant*  $5s(k=0)$  process. These numbers are compatible with the  $\sim 0.2$ -eV experimental location of this peak, particularly if our spectrometer's resolution and laser bandwidth are taken into account. Additionally, the corresponding angular distribution [see inset of Fig. 4(a)] reveals a pronounced *d*-wave contribution, implying a  $5s6p \rightarrow 5s\epsilon d$  ionization step at the four-photon level. Remarkably, the presence of the  $\sim 0.2$ -eV peak persists well above the channel-closing intensity for the nonresonant  $5s(0)$  process [ $\approx 8.6$  TW cm<sup>-2</sup>, as predicted by Eq. (1) for  $\Delta E_{5s^2} = 0$ ]. Its persistence may be attributed to the spatiotemporal intensity distribution of the laser beam and particularly to a laser-atom interaction volume effect [21]. Specifically, it may be attributed to the existence of a “slice” of this volume where the intensity ranges between the resonance threshold and the  $5s(0)$  channel-closing intensities, despite the much higher values the peak intensity may acquire at the laser beam center. Furthermore, the fraction of this slice over the total volume gets smaller with rising intensity, a fact that is reflected in the gradually decreasing magnitude of this peak. On the other hand, even under these conditions Fig. 4(b) (where  $I = 14$  TW cm<sup>-2</sup>) reveals the clear presence of the  $k = 9$  member of the  $\approx 0.2 + k\omega$  eV ( $k \geq 0$ ) *resonant* atomic ATI series, implying that the energy absorbed by the single-active-electron largely exceeds the double-ionization limit of Sr.

The two remaining ATI series should stem from the ionic system. Recall first that the atomic ATI process presented above leads to the production of ground-state Sr<sup>+</sup> ions. Additionally, within this intensity range the data show no evidence of excited ionic-state populations. Hence, the  $5s$  ionic ground state is expected to be the only initial state available for double ionization. This is largely confirmed by the fact that the members of the stronger ionic ATI series (appearing as weak shoulders on the strong atomic ATI peaks) best fit the Sr<sup>2+</sup> ( $5s, k \geq 0$ ) process, observed at  $\approx 1.36 + k\omega$  eV, i.e., very close to the predictions of Eq. (11). They do not shift with intensity, pointing again towards the existence of Freeman resonances. By inspecting the energy-levels diagram of the ion [47], one indeed finds an appreciable number of closely spaced, seven-photon resonant  $nf$  Rydberg states of Sr<sup>+</sup> (see Fig. 1), where the 1.36-eV energy of the  $k = 0$  peak corresponds to  $n \approx 17$ . The relevant threshold intensities of appearance are very low and these Freeman resonances can come into play as soon as MPI of ground-state ions is feasible. In fact, the situation is even more complicated by the additional presence of the quasis resonant  $7d$  state in the vicinity of the sixth photon. Without taking into account any AC Stark shifts this resonance would produce  $\approx 1.32$  eV electrons. However, an accurate estimation of the AC Stark shift due to this double- (or even multiple-) resonant  $7d$ - $nf$  effect should probably necessitate nonperturbative theoretical calculations [7,58,59]. Thus, the behavior of the  $7d$  state is difficult to predict but we may speculate that it could lead to another Freeman resonance and the creation of an electron peak at a different and largely displaced location with respect to the one at 1.36 eV. In fact, this is a possible origin of the third quite weak ionic ATI series at  $\approx 0.78 + k\omega$  eV ( $k = 0,1$ ) observed noticeably in Fig. 4(a), for which there is no other obvious scenario involving the Sr<sup>+</sup> ground state.

## 2. Intensity range above the $\text{Sr}^+$ saturation intensity: Onset of tunneling ionization

A typical PES near the highest available intensity is given in Fig. 4(c). We may observe that as  $I$  increases the higher members of all aforementioned ATI series are practically washed out in favor of a considerable background. The few remaining lower-energy ATI members broaden and their contrast relative to the background decreases with increasing intensity. In fact, the background starts to develop progressively from lower intensities and it is already quite apparent in Fig. 4(b). Its origin, along with the disappearance of the ATI structure, is typically associated with the departure from the MPI picture and the gradual dominance of the tunneling and over-the-barrier ones [15,60]. It is highly probable that the simultaneous presence of ATI and tunneling (and/or over-the-barrier) signatures is to be attributed to the spatiotemporal intensity distribution of the laser beam. It is interesting to note, however, that in our case a large fraction of Sr atoms and/or  $\text{Sr}^+$  ions seem to actually survive ionization until the maximum laser-pulse intensity is reached. On the other hand, the data of Fig. 4(c) were recorded at  $I > I_{\text{OBI},\eta}(\text{Sr})$  and  $I \approx I_{\text{OBI},\eta}(\text{Sr}^+)$ , while  $\gamma > 1$ , a situation reminiscent of strong-field nonlinear ionization of the alkali atoms. In an attempt to resolve this apparent conflict, we adopt the nowadays widely accepted view that the Keldysh parameter alone is not a sufficient indicator for characterizing the ionization regime. Quite frequently, two additional dimensionless indicators are also introduced, namely  $z = 2/\gamma^2$  and  $z_1 = U_p/\omega$  [61,62]. Then, “pure” MPI necessitates  $\gamma > 1$  and simultaneously  $z < 1$  and  $z_1 < 1$  [62]. In the opposite case ( $\gamma < 1$  and simultaneously  $z > 1$  and  $z_1 > 1$ ) tunneling or even over-the-barrier ionization prevails. Accordingly, the regime where  $\gamma > 1$  but either one of  $z$  or  $z_1$  or both of them are larger than unity may be termed as the “intermediate regime,” implying a certain mixture between MPI and tunneling conditions. In our case, starting with  $z_1$ , which is independent of the target system and characterizes solely the laser field, we find that it becomes larger than unity for  $I \geq 26 \text{ TW cm}^{-2}$ , this value being very close to our highest available intensities. Furthermore, one finds  $z(\text{Sr}) \geq 1$  for the comparable lower bound  $I \geq 24 \text{ TW cm}^{-2}$ , while the condition  $z(\text{Sr}^+) \geq 1$  is fulfilled for higher intensities outside the range of the experiment. Evidently, these estimates are fully coherent with respect to the experimental data and point with enough certainty towards the intermediate regime. In fact, this is true for both the atom and the ion (due to  $z_1$  and because  $\text{Sr}^+$  is “fed” by the neutral atom ionization). The above discussion closes our parenthesis on the mixed MPI-tunneling ionization conditions.

## 3. Intensity range below the $\text{Sr}^+$ saturation intensity: CI-ATI signatures

We now turn to the data of Fig. 5 concerning the 3.7–5 –  $\text{TW cm}^{-2}$  intensity range and characterized by the presence of CI-ATI processes. The figure also includes a “reference” PES recorded at  $I \approx 5 \text{ TW cm}^{-2}$ . This latter PES exhibits all the ATI structures discussed above, the difference with respect to the higher-intensity data of Fig. 4 being that most ATI peaks appear now to redshift with increasing intensity. Thus, they reflect primarily the nonresonant  $5s(k)$  and  $\text{Sr}^{2+}(5s, k)$

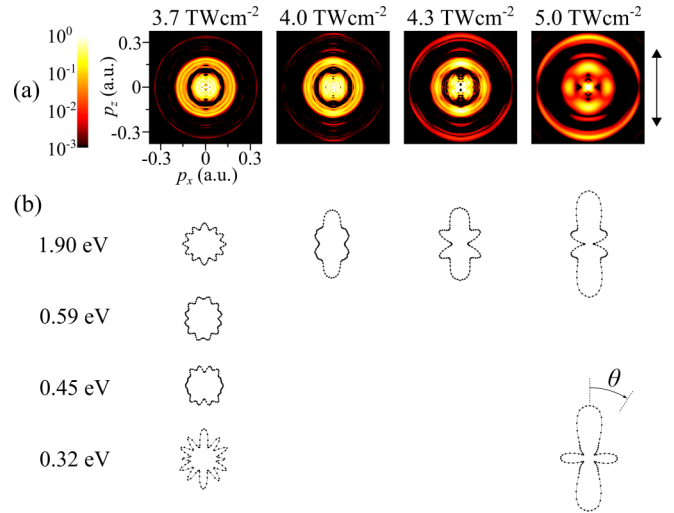


FIG. 6. (a) Inverted (and symmetrized) VMI images recorded at the indicated laser intensities. A logarithmic color scale covering three orders of magnitude is employed for better visibility of their weak features (see the given color bar). The scales of the vertical and horizontal axes are given for a single image but apply to all of them. They are expressed in atomic units of momentum. (b) Normalized polar plots of the PADs obtained from the images in (a) for the indicated electron energies. The evolution of their shapes with intensity and energy is discussed in the text. The angle  $\theta$  with respect to which the PADs are described is shown in one distribution and it is measured with respect to the linear laser polarization. The direction of the latter is denoted by the black double-headed arrow drawn in (a).

processes and they are not yet fully dominated by Freeman resonances. The predictions of Eq. (1) with  $\Delta E_{5s^2} = 0$  are superimposed on the spectrum and reproduce well the locations of the  $5s(1)$  and  $5s(2)$  peaks, but only approximately the location of the  $5s(0)$  one. Using the calculated value for  $\Delta E_{5s^2}$  [57] the agreement worsens for all three peaks. We may conclude that in our case atomic AC Stark shifts are generally far more complicated than that implied by Eq. (1) [59]. Therefore, we set  $\Delta E_{5s^2} = 0$  in the following and assume that the predictions of Eq. (1), although quite useful, indicative, and informative, are quantitatively somewhat inaccurate.

Interestingly, the spectra of Fig. 5 probe weak double-ionization processes within the  $4.3 \leq I \leq 5 \text{ TW cm}^{-2}$  intensity range where no  $\text{Sr}^{2+}$  signal appears in the log-log plot of Fig. 3. This is not fully understood. A possible explanation could involve spatial nonuniformities in the detection efficiency of the MCPs for electrons and ions. Whatever the reason, however, this observation suggests the assumption that no double ionization occurs below the saturation intensity of the  $\text{Sr}^+$  yield should be taken with caution in the following analysis, where, perhaps, a possible weak  $\text{Sr}^{2+}$  production (<5% of the total ion signal as visible on Fig. 3) could be taken into account.

Finally, in order to compare with the lower-intensity data, it is interesting to inspect the  $I \approx 5 \text{ TW cm}^{-2}$  reference PADs given in Fig. 6 for the strong  $5s(k)$  ATI peaks. The PAD of the  $5s(0)$  peak at  $\approx 0.3 \text{ eV}$  is to be compared with the one seen in the inset of Fig. 4(a). They both clearly point towards a



dominant contribution from the  $5s\epsilon d$  continuum (note the strong signal perpendicularly to the laser polarization direction, i.e., at  $\theta = 90^\circ$ ). The  $I \approx 5\text{-TW cm}^{-2}$   $5s(0)$  PAD additionally exhibits a weak sidelobe within the  $0^\circ < \theta < 90^\circ$  interval, probing a quite small  $5s\epsilon g$  contribution. These PAD shapes are similar to the ones reported earlier for four-photon ionization via long ( $\sim\text{ns}$ ) pulses, for either Mg [9] or Sr [12] out of their respective ground states. Based on them one can easily understand the quite asymmetric shape of the  $5s(1)$  PAD at  $\sim 1.9\text{ eV}$  which, as Fig. 6 shows, is characterized by a quasiszero signal at  $\theta = 90^\circ$  and an intense sidelobe within  $0^\circ < \theta < 90^\circ$ , revealing a dominant  $5s\epsilon'f$  contribution. As for the  $5s(2)$  PAD, it is not shown in Fig. 6 but it is identical to the one observed in the inset of Fig. 4(a) and displays a reappearing  $\theta = 90^\circ$  local maximum and a dominant  $5s\epsilon''g$  contribution.

Having presented above all the required reference information, consider next the lower-intensity PES of Fig. 5 which reveal a richer and somewhat more complicated structure with respect to their highest-intensity counterparts. This is also qualitatively evident from the inverted images of Fig. 6. Let us begin the analysis of this different structure by the  $<0.2\text{ eV}$  electron peak which, as laser intensity increases, gradually shifts towards lower energy, weakens in magnitude, and disappears. From Eqs. (1) and (3)–(9) we expect contributions to this signal from the  $4d(0)$  and  $7s(0)$  processes. Nevertheless, one may reasonably expect the  $4d(0)$  ionization pathway to be the by far dominant one due to its much lower number of photons involved (i.e., the much lower nonlinearity of ionization) and the somewhat higher energy of the released electron. The disappearance of this peak is attributed to the closing of this channel ( $\epsilon_{4d,5} = 0$ ) and occurs at  $\approx 4.8\text{ TW cm}^{-2}$  [for  $\Delta E_{5s^2} = 0$  Eq. (1) predicts a somewhat lower disappearance intensity of  $\sim 4\text{ TW cm}^{-2}$ ]. More importantly, its presence ensures the emergence of population in the  $4d$  ionic state, thus signaling the presence of CI-ATI processes. On the other hand, it is verified that double ionization out of this ionic level does not occur since the relevant electron peaks are absent. Finally, the low-intensity spectra of Fig. 5 suggest the weak presence of the  $4d(1)$  peak at  $\approx 1.5\text{ eV}$ , most probably blended with peaks corresponding to other processes discussed below.

Let us deal next with the strongest in magnitude and fairly complex structure observed in the PES of Fig. 5 within the  $0.2\text{--}0.7\text{-eV}$  interval. For low-laser intensity this structure consists of three distinct but partially overlapping lines. As intensity grows it gradually evolves to a single peak clearly attributed to the  $5s(0)$  process. The predictions of Eq. (1) for  $\Delta E_{5s^2} = 0$  drawn in Fig. 5 help us identify the possible candidates for these three peaks at  $I \approx 3.7\text{ TW cm}^{-2}$ . These are the  $5s(0)$  one, the two  $5p_j(0)$  spin-orbit components, and finally, the  $6s(0)$  and  $4f(0)$  ones (the latter two being practically indistinguishable at this energy scale). Among the pathways leading to excited ion formation, the presence of the  $5p_j(0)$  one appears to be the most probable one, due to its much lower nonlinearity of ionization. In fact, since all three peaks are separated by an energy difference of  $\approx 0.1\text{ eV}$  which matches the ionic  $5p$ -level spin-orbit splitting, it is possible that two out of the three peaks belong to the  $5p_{1/2,3/2}(0)$  components. The third peak of the triplet could

just be the  $5s(0)$  one. However, none of the three corresponding PADs of Fig. 6 at  $I \approx 3.7\text{ TW cm}^{-2}$  resembles the one at  $I \approx 5\text{ TW cm}^{-2}$  discussed above and they simply cannot be reproduced by a fit including solely the  $5s\epsilon\ell$ ,  $\ell = 0, 2, 4$  continua. Instead, they all are fairly complex, having several local extrema between  $\theta = 0^\circ$  and  $\theta = 90^\circ$ , they are “symmetric” (approximately equal signals at  $\theta = 0^\circ$  and  $\theta = 90^\circ$ ), and they seem compatible with decay channels corresponding to excited ionic levels, like the  $5p_j\epsilon\ell$  ones. Therefore, at low intensity no peak of the triplet appears to primarily belong to the  $5s(0)$  process. This fact is reminiscent of the observations made in earlier Sr ionization studies with long ( $\sim\text{ns}$ ) and low-intensity pulses, same orders of nonlinearity, and similar wavelength [12], where the excited  $4d$  and  $5p$  ionic levels were found to be by far more populated than the  $5s$  ground state. As for the remaining candidates, namely the  $6s(0)$  and  $4f(0)$  processes, one would expect them to have significantly lower signal than the  $5p_j(0)$  one, due to their much higher nonlinearity and the anticipated substantial atomic population decay to electron-Sr<sup>+</sup> pairs at each photon absorbed within the continuum. Consequently, a more clear picture for the  $6s(0)$  and (to a lesser degree) the  $4f(0)$  channels may be achieved only after considering the origin of the remaining electron peaks.

As seen in Fig. 5, the electron energy expected for the  $5p_j(1)$  process lies very close and overlaps with the  $5s(1)$  one, particularly if the effective laser bandwidth is taken into account. For distinguishing between the two processes and observing their competition with varying intensity we invoke the PAD at the maximum of the relevant peak located at  $\approx 1.9\text{ eV}$ . The intensity evolution of this PAD is given in Fig. 6, along with the high-intensity reference probing the  $5s(1)$  pathway already discussed above. With respect to that PAD, the  $I \approx 3.7\text{ TW cm}^{-2}$  one is completely different and evidently related to the PADs of the triplet structure within the  $0.2\text{--}0.7\text{-eV}$  range for the same intensity. Specifically, this PAD also exhibits several local extrema between  $\theta = 0^\circ$  and  $\theta = 90^\circ$  and comparable signals at these two angles. Consequently, this PAD excludes the  $5s(1)$  pathway and, more notably, it probes the  $5p_j(1)$  process, because there are no other (single or double-ionization) processes expected at this energy. Figure 6 shows that as intensity increases the  $5s(1)$  and  $5p_j(1)$  pathways coexist, resulting in an incoherent sum of two PADs [63] and a  $\theta = 90^\circ$  signal gradually diminishing and favoring the pure  $5s(1)$  PAD. As a last remark on this spectral feature, we may anticipate that the emergence of a single broad  $5p_j(1)$  peak instead of a  $j = 1/2, 3/2$  doublet is to be most probably attributed to its blend with the peaks of other processes and to the increased effective laser bandwidth.

There is a last remaining quite weak peak to be discussed. It is observed in the low-intensity PES of Fig. 5 at  $\approx 1.0\text{ eV}$  and disappears for  $I > 4.3\text{ TW cm}^{-2}$ . Given the proven presence of the  $5p_j(1)$  pathway, the energy location and low magnitude of this peak correlates well with the  $6p(0)$  process [and would also support a contribution of the  $6s(0)$  channel related to the  $\approx 0.59\text{-eV}$  peak of the  $0.2\text{--}0.7\text{-eV}$  triplet]. However, although unlikely, this peak could also emerge from the Sr<sup>2+</sup>( $5p_j, 0$ ) double-ionization process. If only the ponderomotive shift is included in Eq. (12) we indeed expect an  $\sim 1\text{-eV}$  energy peak, while if the calculated AC Stark shift of the  $5p_j$  levels [56] is

also included its energy is found to be  $\ll 1$  eV. Still, an  $\sim 1$ -eV peak may be produced if Freeman resonances are considered for the *resonant*  $\text{Sr}^{2+}(5p_j, 0)$  pathway, but such a possibility is rather weak because the observed peak slightly downshifts with increasing intensity. The same reasoning applies also for the double-ionization pathway out of the  $6s$  ionic level.

The findings presented above reveal the presence of CI-ATI processes at low intensity. Indeed, the analysis of relevant PES and PADs led to the detection of, at least, the  $4d(0)$  and  $5p_j(1)$  CI-ATI pathways (for the former the other less probable candidate is also a CI-ATI process). Moreover, the emergence of channel  $5p_j(1)$  (which implies the absorption of three CI-ATI photons) requires the presence of the  $5p_j(0)$  one as a prerequisite. For the latter pathway the relevant electron peak appears to be indeed present in the PES, but it is blended with other peaks within the  $0.2$ – $0.7$ -eV range. This blend renders the detailed assignment of these peaks rather incomplete and difficult to achieve at present. Nevertheless, it is to be emphasized that our study does not mainly focus on their identification. Instead, it focuses on the proved observation of CI-mediated absorption processes, as well as on the transition towards the single-active-electron behavior as intensity increases. These facts are very clearly registered in Figs. 5 and 6. It is finally important to note that the proof of CI-ATI presence is here based solely on experimental data and without any prior or complementary reference to theoretical calculations. On the other hand, theoretical modeling would be, of course, highly desirable for providing further insight.

## V. CONCLUDING REMARKS AND FUTURE PERSPECTIVES

We have presented measurements of a photoelectron spectroscopy study devoted to the nonlinear single and double ionization of Sr atoms interacting with  $\approx 800$ -nm laser pulses of  $\approx 25$ -fs duration. Within the examined laser-intensity range, photoelectron-energy spectra and angular distributions probe two consecutive transitions between processes of qualitatively different nature. At low intensity,  $I \leq 4 \text{ TW cm}^{-2}$ , the recorded data revealed the production of excited singly charged ions, thus proving that single ionization proceeds via excitation ladders composed by quasis resonant doubly excited states. The emergence of dense manifolds of the latter states just above (or even below) the first ionization threshold characterizes Sr and the other alkaline-earth-metal atoms and it has been the point of focus in the present work because of the strong interaction between the two valence electrons it implies. However, this mechanism is rather abruptly turned off at higher intensity where it is replaced by a single-active-electron one which is probed by the observation of typical atomic and ionic ATI structures. Finally, by increasing the intensity even further the MPI picture gradually diminishes in favor of tunneling ionization.

We have shown that this last MPI-to-tunneling ionization transition, while unexpected on the basis of the Keldysh parameter alone, can be explained by additionally implicating several other indicators for properly characterizing the ionization regime. These indicators provided an estimated intensity of  $\approx 24 \text{ TW cm}^{-2}$  that separates the two regimes and this value roughly agrees with experimental observations. On the

contrary, there is no clear explanation concerning the origin of the sharp switching off of the electron-correlation mediated processes. In the relevant earlier work devoted to Mg atom [32], the results of a rate equation model attributed this switching off to the small population accumulated to the excited ionic state as compared to the ground state of  $\text{Mg}^+$ , as well as to the fast depletion of this excited population with increasing intensity. This does not seem to be the case in the present experiment, where excited ionic-state populations appear to be considerably larger than the ionic ground-state one at low intensity. Thus, an alternative explanation is required. In the absence of any theoretical results stemming from an appropriate model including two-electron effects, we tentatively attribute the effect to the blueshift of the  $4d5p\ ^1F_3$  and  $5s7p\ ^1P_1$  three-photon near-resonant states with increasing intensity. These states exhibit considerable mixing among doubly excited and singly excited (Rydberg) configurations and constitute the first ladder step. Thus, their blueshift (either the ponderomotive or a more subtle AC Stark one) would push them out of resonance, destroying this way the whole ladder structure. In addition, the  $4d(0)$  channel closing that occurs within the same intensity range may also create unfavorable CI-ATI conditions.

Interestingly, both of the above assumptions can be experimentally put to the test, since they can be both remedied, in principle, by increasing the photon energy. An obvious solution would be to repeat the earlier experiments in Sr performed at  $\approx 725 \pm 10 \text{ nm}$  [12], by replacing the long ( $\sim$ ns) pulses with ultrashort ( $\sim$ fs) ones. These wavelengths can be easily obtained nowadays by optical parametric amplifier systems. This choice, however, presents several shortcomings. First, there is a considerable overlap between the electron energies produced by different pathways. Second, the aforementioned resonances are expected to resonantly upshift at rather high intensities, the latter largely exceeding the  $\text{Sr}^+$  yield saturation intensity which is expected to be close to the presently observed one. Therefore, the radiation wavelength needs to be more judiciously chosen. Assuming only the presence of  $U_p$ , we estimate that an interesting choice would be to employ  $\approx 740$ – $745$ -nm radiation. The latter range ensures the  $4d(0)$  channel closing would correspond to an intensity of  $\sim 10 \text{ TW cm}^{-2}$ , which is much higher than the one witnessed at the present experiment and close to the saturation intensity of  $\text{Sr}^+$ . More importantly, at about the same intensity the upshifted aforementioned states would now be brought into three-photon resonance. Finally, most ionization pathways are expected to produce fairly well-resolved electron peaks.

The aim of the above proposals is to push the observation of CI-ATI processes at intensities on the order of  $10 \text{ TW cm}^{-2}$  or higher. Under such conditions we may even envision an increasing probability to additionally observe CI-mediated direct (or nonsequential) double ionization within the MPI regime. If successful, such experiments would allow bridging with observations of nonsequential double-ion production within the tunneling regime [34,37–39]. In this respect, it is evident that the presence of sequential double-ionization processes would play an obscuring role. Consequently, the employment of coincidence studies [64] at some stage of this research direction would be unavoidable. Furthermore, it would be highly desirable for the choice of the relevant

experimental parameter space to be guided by two-active-electron theoretical modeling (see, for example, Refs. [65,66]). Therefore, recognizing that present-day experimental and (to some extent) theoretical techniques have

reached a certain degree of maturity, we hope that the work presented here may initiate a different, albeit promising, pathway towards the study of electron-correlation effects in the ultrafast regime.

- [1] I. S. Aleksakhin, I. P. Zapesochnyi, and V. V. Suran, *JETP Lett.*, **26**, 11 (1977); I. S. Aleksakhin, N. B. Delone, I. P. Zapesochnyi, and V. V. Suran, *Sov. Phys. JETP* **49**, 447 (1979).
- [2] D. Feldmann, H. J. Krautwald, and K. H. Welge, *J. Phys. B: At. Mol. Phys.* **15**, L529 (1982); D. Feldmann, H. J. Krautwald, S. L. Chin, A. von Hellfeld, and K. H. Welge, *ibid.* **15**, 1651 (1982); N. B. Delone, V. V. Suran, and B. A. Zon, *Multiphoton Ionization of Atoms*, edited by S. L. Chin and P. Lambropoulos, (Academic, New York, 1984); J. L. Dexter, S. M. Jaffe, and T. F. Gallagher, *J. Phys. B: At. Mol. Phys.* **18**, L735 (1985); G. Petite, and P. Agostini, *J. Phys. (Paris)* **47**, 795 (1986); P. Lambropoulos, X. Tang, P. Agostini, G. Petite, and A. L'Huillier, *Phys. Rev. A*, **38**, 6165 (1988).
- [3] P. Camus, M. Kompitsas, S. Cohen, C. A. Nicolaides, M. Aymar, M. Crance, and P. Pillet, *J. Phys. B: At. Mol. Phys.* **22**, 445 (1989); Y. Zhu, R. R. Jones, W. Sandner, T. F. Gallagher, P. Camus, P. Pillet, and J. Boulmer, *ibid.* **22**, 585 (1989); T. T. Bernat, I. I. Bondar, and V. V. Suran, *Opt. Spectrosc.* **71**, 22 (1991); S. G. Nakhate, S. A. Ahmad, M. A. N. Razvi, and G. D. Saksena, *J. Phys. B* **24**, 4973 (1991); D. A. Tate, D. G. Papaioannou, and T. F. Gallagher, *ibid.* **24**, 1953 (1991); D. G. Papaioannou, D. A. Tate, and T. F. Gallagher, *ibid.* **25**, 2517 (1992); H. K. Haugen and H. Stapelfeldt, *Phys. Rev. A* **45**, 1847 (1992); N. J. van Druten and R. Trainham, *ibid.* **50**, 1593 (1994); I. I. Bondar, and V. V. Suran, *JETP Lett.* **68**, 837 (1998).
- [4] D. Kim, S. Fournier, M. Saeed, and L. F. DiMauro, *Phys. Rev. A*, **41**, 4966 (1990).
- [5] P. Lambropoulos, P. Maragakis, and J. Zhang, *Phys. Rep.* **305**, 205 (1998), and references therein.
- [6] I. I. Bondar, V. V. Suran, and M. I. Dudich, *J. Phys. B: At. Mol. Phys.* **33**, 4243 (2000); I. I. Bondar and V. V. Suran, *ibid.* **35**, 3391 (2002).
- [7] I. Lontos, A. Bolovinos, S. Cohen, and A. Lyras, *Phys. Rev. A*, **70**, 033403 (2004); I. Lontos, S. Cohen, and A. Lyras, *J. Phys. B: At. Mol. Phys.* **43**, 095602 (2010).
- [8] D. I. Bondar, G. L. Yudin, W. K. Liu, M. Y. Ivanov, A. D. Bandrauk, *Phys. Rev. A*, **83**, 013420 (2011); I. I. Bondar, V. V. Suran, and D. I. Bondar, *ibid.* **88**, 023407 (2013).
- [9] A. Dimitriou, S. Cohen, and A. Lyras, *J. Phys. B: At. Mol. Phys.* **44**, 135001 (2011); A. Dimitriou, S. Cohen, A. Lyras, and I. Lontos, *ibid.* **45**, 205003 (2012).
- [10] I. Lontos, A. Bolovinos, and S. Cohen, *J. Phys. B: At. Mol. Phys.* **41**, 045601 (2008).
- [11] A. Dimitriou and S. Cohen, *Phys. Rev. A*, **90**, 012513 (2014).
- [12] A. Dimitriou and S. Cohen, *Eur. Phys. J. D* **69**, 238 (2015).
- [13] *Atoms in Intense Laser Fields*, edited by M. Gavrilu, *Advances in Atomic, Molecular and Optical Physics, Suppl. 1* (Academic, New York, 1992).
- [14] L. V. Keldysh, *Sov. Phys. JETP* **20**, 1307 (1965).
- [15] N. B. Delone and V. P. Krainov, *Phys. Usp.* **41**, 469 (1998).
- [16] P. Agostini, F. Fabre, G. Mainfray, G. Petite, and N. K. Rahman, *Phys. Rev. Lett.* **42**, 1127 (1979).
- [17] M. V. Ammosov, N. B. Delone, M. Yu. Ivanov, I. I. Bondar, and A. V. Masalov, *Adv. At., Mol., Opt. Phys.* **29**, 33 (1991); G. N. Gibson, R. R. Freeman, T. J. McIlrath, and H. G. Muller, *Phys. Rev. A*, **49**, 3870 (1994).
- [18] K. J. Schafer, B. Yang, L. F. DiMauro, and K. C. Kulander, *Phys. Rev. Lett.* **70**, 1599 (1993).
- [19] M. Schuricke, G. Zhu, J. Steinmann, K. Simeonidis, I. Ivanov, A. Kheifets, A. N. Grum-Grzhimailo, K. Bartschat, A. Dorn, and J. Ullrich, *Phys. Rev. A*, **83**, 023413 (2011).
- [20] T. Morishita and C. D. Lin, *Phys. Rev. A*, **87**, 063405 (2013).
- [21] S.-D. Jheng and T. F. Jiang, *J. Phys. B: At. Mol. Phys.* **46**, 115601 (2013).
- [22] N. A. Hart, J. Strohaber, A. A. Kolomenskii, G. G. Paulus, D. Bauer, and H. A. Schuessler, *Phys. Rev. A*, **93**, 063426 (2016).
- [23] J. Yuan, S. Liu, X. Wang, Z. Shen, Y. Ma, H. Ma, Q. Meng, T.-M. Yan, Y. Zhang, A. Dorn, M. Weidemüller, D. Ye, and Y. Jiang, *Phys. Rev. A*, **102**, 043112 (2020).
- [24] D. Zille, D. Adolph, S. Skruszewicz, A. M. Sayler, and G. G. Paulus, *New J. Phys.* **22**, 083021 (2020).
- [25] P. Wessels, B. Ruff, T. Kroker, A. K. Kazansky, N. M. Kabachnik, K. Sengstock, M. Drescher, and J. Simonet, *Commun. Phys.* **1**, 32 (2018).
- [26] A. Bunjac, D. B. Popovic, and N. S. Simonovic, *Phys. Lett. A* **394**, 127197 (2021).
- [27] F. Morales, M. Richter, S. Patchkovskii, and O. Smirnova, *Proc. Natl. Acad. Sci.* **108**, 16906 (2011).
- [28] M. Aymar, C. H. Greene, and E. Luc-Koenig, *Rev. Mod. Phys.* **68**, 1015 (1996).
- [29] M. Sukharev, E. Charron, and A. Suzor-Weiner, *Phys. Rev. A*, **66**, 053407 (2002).
- [30] J. Zhang and P. Lambropoulos, *Phys. Rev. Lett.* **77**, 2186 (1996).
- [31] L. Nikolopoulos, G. Buica-Zloh, and P. Lambropoulos, *Eur. Phys. J. D* **26**, 245 (2003).
- [32] D. Xenakis, N. E. Karapanagioti, D. Charalambidis, H. Bachau, and E. Cormier, *Phys. Rev. A*, **60**, 3916 (1999).
- [33] L. D. Van Woerkom, M. J. Nandor, M. A. Walker, G. D. Gillen, and H. G. Muller, *Laser Phys.* **11**, 982 (2001).
- [34] G. D. Gillen, M. A. Walker, and L. D. Van Woerkom, *Phys. Rev. A*, **64**, 043413 (2001).
- [35] G. D. Gillen and L. D. Van Woerkom, *Phys. Rev. A*, **68**, 033401 (2003).
- [36] E. Papastathopoulos, M. Strehle, and G. Gerber, *Chem. Phys. Lett.* **408**, 65 (2005).
- [37] H. Kang, S. Chen, Y. L. Wang, W. Chu, J. P. Yao, J. Chen, X. J. Liu, Y. Cheng, and Z. Z. Xu, *Phys. Rev. A*, **100**, 033403 (2019).
- [38] Y. H. Lai, X. Wang, Y. Li, X. Gong, B. K. Talbert, C. I. Blaga, P. Agostini, and L. F. DiMauro, *Phys. Rev. A*, **101**, 013405 (2020).
- [39] H. P. Kang, S. Chen, W. Chu, J. P. Yao, J. Chen, X. J. Liu, Y. Cheng, and Z. Z. Xu, *Opt. Express* **28**, 19325 (2020).
- [40] A. T. J. B. Eppink and D. H. Parker, *Rev. Sci. Instrum.* **68**, 3477 (1997).

- [41] M. M. Harb, S. Cohen, E. Papalazarou, F. Lépine, and C. Bordas, *Rev. Sci. Instrum.* **81**, 125111 (2010).
- [42] M. J. J. Vrakking, *Rev. Sci. Instrum.* **72**, 4084 (2001).
- [43] K. Zhao, T. Colvin, W. T. Hill, and G. Zhang, *Rev. Sci. Instrum.* **73**, 3044 (2002); G. M. Roberts, J. L. Nixon, J. Lecointre, E. Wrede, and J. R. R. Verlet, *ibid.* **80**, 053104 (2009).
- [44] A. Talebpour, C.-Y. Chien, Y. Liang, S. Larochelle, and S. L. Chin, *J. Phys. B: At. Mol. Phys.* **30**, 1721 (1997); C. Guo, M. Li, J. P. Nibarger, and G. N. Gibson, *Phys. Rev. A* **58**, R4271(R) (1998).
- [45] W. Becker, E. Grasbon, R. Kopold, D. B. Milošević, G. G. Paulus, and H. Walther, *Adv. At., Mol. Opt. Phys.* **48**, 35 (2002).
- [46] R. R. Freeman, P. H. Bucksbaum, H. Milchberg, S. Darack, D. Schumacher, and M. E. Geusic, *Phys. Rev. Lett.* **59**, 1092 (1987).
- [47] Y. Ralchenko, A. Kramida, and J. Reader, and NIST ASD Team, 2011 NIST Atomic Spectra Database (version 4.1), <http://physics.nist.gov/asd>
- [48] J. E. Sansonetti and G. Nave, *J. Phys. Chem. Ref. Data* **39**, 033103 (2010).
- [49] M. Wollenhaupt, M. Krug, J. Köhler, T. Bayer, C. Sarpe-Tudoran, and T. Baumert, *Appl. Phys. B* **95**, 245 (2009).
- [50] C. L. Vaillant, M. P. A. Jones, and R. M. Potvliege, *J. Phys. B: At. Mol. Phys.* **47**, 155001 (2014).
- [51] E. Luc-Koenig, Aymar M, J.-M. Lecomte, and A. Lyras, *J. Phys. B: At. Mol. Phys.* **31**, 727 (1998).
- [52] C. C. Chu, H. S. Fung, H. H. Wu, and T. S. Yih, *J. Phys. B: At. Mol. Phys.* **31**, 3843 (1998), and references therein.
- [53] C. J. Dai and J. Lu, *J. Phys. B: At. Mol. Phys.* **29**, 2473 (1996).
- [54] M. Aymar, J.-M. Lecomte, C. C. Chu, H. S. Fung, H. H. Wu, and T. S. Yih, *J. Phys. B: At. Mol. Phys.* **31**, 5135 (1998).
- [55] L. Hamonou, T. Morishita, and O. I. Tolstikhin, *Phys. Rev. A*, **86**, 013412 (2012).
- [56] J. Kaur, S. Singh, B. Arora, and B. K. Sahoo, *Phys. Rev. A* **92**, 031402(R) (2015).
- [57] S. G. Porsev, A. D. Ludlow, M. M. Boyd, and J. Ye, *Phys. Rev. A* **78**, 032508 (2008); K. Guo, G. Wang, and A. Ye, *J. Phys. B: At. Mol. Phys.* **43**, 135004 (2010).
- [58] S. A. Kotochigova, *J. Opt. Soc. Am. B* **9**, 1215 (1992).
- [59] R. M. Potvliege and S. Vučić, *Phys. Scr.* **74**, C55 (2006).
- [60] E. Mevel, P. Breger, R. Trainham, G. Petite, and P. Agostini, *Phys. Rev. Lett.* **70**, 406 (1993).
- [61] H. R. Reiss, *Phys. Rev. Lett.* **101**, 043002 (2008); **101**, 159901(E) (2008).
- [62] K. D. Schultz, C. I. Blaga, R. Chirla, P. Colosimo, J. Cryan, A. M. March, C. Roedig, E. Sistrunk, J. Tate, J. Wheeler, P. Agostini, and L. F. DiMauro, *J. Mod. Opt.* **54**, 1075 (2007); P. Agostini, and L. F. DiMauro, *Adv. At. Mol. Opt. Phys.* **61**, 117 (2012); L. F. DiMauro, and C. A. Roedig, *Progress in Ultrafast Intense Laser Science, X, Vol. 106*, Springer Series in Chemical Physics (Springer, 2014), Chap. 5, pp. 65–76.
- [63] M. Goto and K. Hansen, *Phys. Scr.* **86**, 035303 (2012).
- [64] B. Witzel, N. A. Papadogiannis, and D. Charalambidis, *Phys. Rev. Lett.* **85**, 2268 (2000); H. Schmidt-Böcking *et al.*, *Nucl. Phys. A* **737**, 306 (2004).
- [65] K. Amini, J. Biegert, F. Calegari, and A. Chacón, *Rep. Prog. Phys.* **82**, 116001 (2019).
- [66] G. D. Borisova *et al.*, *J. Phys. Commun.* **4**, 055012 (2020).