Asymmetry of the photoelectron momentum distribution from molecular ionization in elliptically polarized laser pulses

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(Received 13 November 2018; published 24 January 2019)

We theoretically study the photoelectron momentum distributions (PMDs) of H$_2^+$ in elliptically polarized laser fields by numerically solving the time-dependent Schrödinger equation. Depending on the internuclear distance of H$_2^+$, the PMDs reveal different asymmetries with respect to the major and minor axes of the laser ellipse. At an internuclear distance corresponding to the enhanced ionization region, the asymmetry of the PMDs is opposite to that of the companion atom. We attribute this phenomenon to the effect of the internal scattered electrons, which are mainly released at an earlier ionization moment with respect to the field maximum within each half cycle. We further show that the asymmetry of the PMDs in the elliptical laser fields can be used to identify the instant when the electron wave packet is released from the molecule.

DOI: 10.1103/PhysRevA.99.013431

I. INTRODUCTION

Atoms or molecules exposed in a strong laser pulse can be tunnel ionized with releasing an electron wave packet. The released electron wave packet can be driven back to the parent ion by the oscillating laser field, which results in many highly nonlinear phenomena, such as high-order above-threshold ionization [1–4], nonsequential double ionization [5,6], and high-order harmonics generation [7–9]. Those processes all depend on the instant when the electron is released within a half cycle of the laser field. For an atom, the bound electron directly tunnels through the suppressed Coulomb barrier. This leads to the expectation that, when the laser field is maximum, the tunneling barrier is the thinnest and the ionization rate of the atom is the highest [10]. In an elliptically polarized laser field, it is expected that the maximal ionization rate corresponds to the instant when the laser field is along the major axis of the laser ellipse and the photoelectron momentum distributions (PMDs) is symmetric with respect to the major and minor axes of the laser ellipse according to the strong-field approximation [11]. In fact, this symmetry is broken by the effect of the Coulomb potential (known as Coulomb asymmetry) [12–14], and the main emission angle in the PMD is shifted to a later ionization moment with respect to the maximum of the laser field.

Compared with the case of an atom, the tunneling ionization of a molecule is more complicated due to the polyatomic Coulomb potential. We take the simplest molecule, i.e., H$_2^+$, as an example. As shown in Fig. 1, there might be two barriers formed by the combination of the H$_2^+$ potential and the external field. When the internuclear distance is short, the electron tunnels through the outer barrier into the continuum, which is similar to an atom. When the internuclear distance increases to a critical value, the inner barrier is formed for the molecule. The electron might tunnel through the thin inner barrier from the up-field core. As a result, the ionization rate increases at this critical internuclear distance, which is known as charge-resonance enhanced ionization (EI) [15–23]. The enhancement of the ionization rate and the tunneling site for different internuclear distances have been theoretically predicted and experimentally demonstrated in molecules. Recently, it has been further shown that the shape of the momentum distribution along the laser propagation direction can be used as an indicator of the tunneling site [23]. Moreover, in the EI region of the molecule, it is predicted that there are multiple ionization bursts [18,19,24] within a half cycle of the laser field because of the transient localization of the electrons at one of the nuclei. This indicates a rather complex electron dynamics on the sub-laser-cycle timescale in molecules. Up to now, it is still unclear when the electron wave packets are mainly released from the molecule for different internuclear distances. Furthermore, the spherical symmetry of the Coulomb potential is broken for molecules, which might have a great effect on the electron dynamics of the Coulomb asymmetry in an elliptically polarized laser pulse [25].

In this paper, we study the PMDs of H$_2^+$ in elliptically polarized laser fields by numerically solving the time-dependent Schrödinger equation (TDSE). The PMD of H$_2^+$ reveals an asymmetry with respect to the major and minor axes of the laser ellipse, which depends on the internuclear distance. For an internuclear distance corresponding to the EI region, we find that the asymmetry of the PMDs is opposite to that of the companion atom. The asymmetry of the PMDs can not only be used to identify the tunneling site of the molecule, but also

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scattered by the down-field core on its subsequent motion. The inset: the internal scattered electron is in a strong electric field. The electron can be released either from the up-field core and then scattered by the down-field core on its subsequent motion.

II. THEORETICAL METHODS

A. Numerical solutions of the TDSE

We solve the two-dimensional TDSE of fixed-nuclei $H_2^+$ in an elliptically polarized laser pulse. The TDSE of $H_2^+$ can be written as [atomic units (a.u.) are used throughout unless specified otherwise]

$$\frac{\partial}{\partial t} \Psi(r, t) = H(r, t) \Psi(r, t),$$

where $\Psi(r, t)$ is the electron wave function, $r = (x, y)$ denotes the electron position, and $H(r, t)$ is the Hamiltonian. In the length gauge, $H(r, t)$ is given by

$$H(r, t) = -\frac{1}{2} \nabla^2 + V(r) + r \cdot E(t),$$

where the Coulomb potential is expressed as

$$V(r) = -\frac{1}{\sqrt{(x-R/2)^2 + y^2 + a}} - \frac{1}{\sqrt{(x + R/2)^2 + y^2 + a}},$$

with $R$ being the internuclear distance, $a$ is the soft-core parameter. For $R = 4$ and 8 a.u., the soft-core parameter is chosen to be 0.72 and 0.723, corresponding to the ionization potential of $H_2^+$ of 0.796 and 0.614 a.u., respectively. The elliptically polarized laser field is given by

$$E(t) = E_0 f(t) \left[ \frac{1}{\sqrt{1 + \epsilon^2}} \cos(\omega t) \hat{e}_x + \frac{\epsilon}{\sqrt{1 + \epsilon^2}} \sin(\omega t) \hat{e}_y \right],$$

where $E_0$ is the electric-field amplitude, $\epsilon$ is the ellipticity, and $\omega$ is the angular frequency of the laser pulse. The laser pulse envelope $f(t) = \sin^2(\pi t / T_p)$ is employed with a duration of $T_p = 13T_0$, where $T_0$ is the period of the 800-nm laser field.

To ensure all the ionized components move away from the core, the wave function is further propagated for an additional two optical cycles at the end of the pulse. We use a splitting-operator fast-Fourier transform method to numerically solve the TDSE of the diatomic molecule [26,27]. The ground state of the molecule is obtained by imaginary-time propagation. The whole wave function at any given time $\tau$ is split into two parts,

$$\Psi(\tau) = \Psi(\tau) [1 - F_1(R_c)] + \Psi(\tau) [F_1(R_c)],$$

$$= \Psi_I(\tau) + \Psi_{II}(\tau).$$

Here, $F_1(R_c) = 1/(1 + e^{-(r-R_c)/\Delta})$ is an absorbing function that separates the whole space into the inner $(0 \rightarrow R_c)$ wave-function $\Psi_I(\tau)$ and outer $(R_c \rightarrow R_{max})$ wave-function $\Psi_{II}(\tau)$. $\Psi_I$ and $\Psi_{II}$ are propagated under the full Hamiltonian numerically and the Volkov Hamiltonian analytically, respectively, $\Delta = 8$ a.u. represents the width of the crossover region. The photoelectron spectra are obtained from the outer wave function. At each time step, $\Psi_{II}$ is transformed into momentum space,

$$C(p, \tau) = \int \Psi_{II}(\tau) e^{-i[p + A(\tau)]r / 2\pi} d^2r,$$

where $A(\tau)$ is the vector potential of the laser field and $\Psi_{II}$ is propagated to the final time as

$$\Psi_{II}(\infty, \tau) = \exp \left\{ -i \int_{\tau}^{\infty} \frac{1}{2} [p + A(\tau')]^2 d\tau' \right\} \times C(p, \tau) e^{ipr / 2\pi} d^2p,$$

so the final momentum distribution is obtained as

$$\frac{dP(p)}{dE} d\theta = \sum_\tau C(p, \tau)^2.$$
B. Classical-trajectory model

The classical-trajectory model is used to analyze the internal scattered electrons (the inset of Fig. 1) at the EI region of H$_2^+$, which is similar to the classical rescattering model in Refs. [28,29]. Briefly, the electron is released at one of the nuclei, i.e., $r = (-4, 0)$ a.u. with zero initial longitudinal momentum along the instantaneous laser field direction. A nonzero initial transverse momentum $v_{0\perp}$ is used to compensate the electron motion induced by the rotating electric field. In addition, the ionization time of the electron is restricted within $t_i \in (0.25, 0.75)t_0$. The electron motion in the laser field is obtained by integrating the classical Newtonian equation without considering the Coulomb effect. Thus, the velocities of the electron along the $x$ and $y$ directions, respectively, can be obtained by

$$v_x(t) = -\int_0^t E_x(t')dt' = -\frac{E_0'}{\omega}[\sin(\omega t) - \sin(\omega t_0)] + v_{x0},$$

$$v_y(t) = -\int_0^t E_y(t')dt' = \frac{E_0'}{\omega}[\cos(\omega t) - \cos(\omega t_0)] + v_{y0},$$

(9)

where $E_0' = \frac{E_0}{\sqrt{1 + \varepsilon^2}}$, $v_{x0} = v_\perp \sin \phi$, $v_{y0} = v_\perp \cos \phi$, and $\phi = \arctan(E_y'/E_x')$. The electron positions $x$ and $y$, respectively, can be written as

$$x(t) = -\int_0^t v_x(t')dt' = -\frac{E_0'}{\omega^2}[\cos(\omega t) - \cos(\omega t_0)] + \omega \sin(\omega t_0)(t - t_0) + v_{x0}(t - t_0) + x_0,$$

$$y(t) = -\int_0^t v_y(t')dt' = \frac{E_0'}{\omega^2}[\sin(\omega t) - \sin(\omega t_0)] - \omega \cos(\omega t_0)(t - t_0) + v_{y0}(t - t_0) + y_0,$$

(10)

where the electron initial positions are $x_0 = -4$ and $y_0 = 0$ a.u. The scattering time $t_s$ is obtained by the conditions $x(t_s) = 4$ and $y(t_s) = 0$ a.u. The velocities of the electron along the $x$ and $y$ directions, respectively, at the moment of scattering can be given by Eq. (9),

$$v_{x_s} = -\frac{E_0'}{\omega}[\sin(\omega t_s) - \sin(\omega t_0)] + v_{x0},$$

$$v_{y_s} = \frac{E_0'}{\omega}[\cos(\omega t_s) - \cos(\omega t_0)] + v_{y0}.$$

(11)

The electron energy $E_e$ and the angle $\phi_e$, respectively, with respect to the $x$ direction at the instant of scattering can be calculated by

$$E_e = \frac{1}{2}(v_{x_s}^2 + v_{y_s}^2),$$

$$\phi_e = \arccos \frac{v_{x_s}}{\sqrt{v_{x_s}^2 + v_{y_s}^2}}.$$

(12)

With considering the zero scattering angle, the final momentum of the internal scattered electron along the $x$ and $y$ directions, respectively, can be obtained by

$$p_x = \sqrt{2E_e}\cos\phi_e + \frac{E_0'}{\omega}\sin(\omega t_s),$$

$$p_y = \sqrt{2E_e}\sin\phi_e - \frac{E_0'}{\omega}\cos(\omega t_s).$$

(13)

III. RESULTS AND DISCUSSIONS

We first show the field-driven momentum $p = -A(t_0)$ in Fig. 2(a) for the elliptically polarized laser field with an ellipticity of 0.4, where $t_0$ is the ionization time for the electron. The electric field of the laser pulse consists of four segments per cycle, i.e., $0 < t < 0.25t_0$ (red), $0.25t_0 < t < 0.5t_0$ (blue), $0.5t_0 < t < 0.75t_0$ (green), and $0.75t_0 < t < t_0$ (black). (b)–(d) show the PMDs of (b) a model atom, (c) H$_2^+$ with $R = 4$ a.u., and (d) H$_2^+$ with $R = 8$ a.u., respectively, in an elliptically polarized laser field ($\varepsilon = 0.4$, $I = 1.5 \times 10^{14}$ W/cm$^2$, and $\lambda = 800$ nm).
angle is much closer to the major axis of the laser ellipse. When the internuclear distance increases to 8 a.u., as shown in Fig. 2(d), the PMD reveals an opposite asymmetry with that of the atom. Most electrons are distributed in the first and third quadrants of the momentum plane, which correspond to the rising edge of the laser field along the major axis of the laser ellipse, i.e., $0.25T_0 < t < 0.75T_0$ and $0.75T_0 < t < T_0$.

To shed light on the dependence of the asymmetry on the internuclear distance, we have traced the time evolution of the electron density along the molecular axis direction by the TDSE as shown in Fig. 3. For the case of the atom [Fig. 3(a)], the electrons are mainly released near the field maxima of $t = 5.5T_0$ and $t = 6T_0$ as indicated by the arrows. For the case of H$_2^+$ at $R = 4$ a.u. [Fig. 3(b)], the bound electron density is transiently localized at one of the two protons at $x = \pm 2$ a.u. within a half cycle of the laser field [18,19]. We note that the electrons are mainly released after the field maxima when the bound electron is localized at the down-field core as indicated by the arrows. According to $p = -A(t_0)$, the main emission angle will shift to the major axis compared with that of the atom, which agrees with the PMD in Fig. 2(c). The case is more complex when the internuclear distance increases to 8 a.u. (EI region). As shown in Fig. 3(c), the electron is released in several bunches within a half cycle of the laser field, which is consistent with previous studies [19]. One can clearly see the internal scattered electrons as indicated by the arrows of A, B, A', and B', which is released from the up-field core and then scattered by the down-field core on its subsequent motion [22]. Notably, those internal scattered electrons are mainly released at the rising edge of the laser field along the major axis of the laser ellipse. According to the mapping relation between the ionization time and the final momentum, we can infer that those internal scattered electrons are mainly distributed in the first and third quadrants of the final momentum plane. This is consistent with the PMD in Fig. 2(d).

To show why the internal scattered electrons are mainly released at the rising edge of the laser field along the major axis of the laser ellipse at the EI regions, we resort to the classical-trajectory model. Figure 4(a) shows the electron energy at the instant of scattering with respect to the ionization time (blue asterisk) and the scattering time (orange cross) for the internal scattered electron in H$_2^+$ with $R = 8$ a.u. calculated by the classical-trajectory model. (b) shows the typical trajectories of the internal scattered electron in the direction of the molecular axis. (c) The absolute value of the initial transverse momentum with respect to the ionization time. In (a)−(c), the gray solid curves are the laser fields along the major axis of the laser ellipse. (d) exhibits the final two-dimensional momenta of the internal scattered electron with assuming zero scattering angle.
of the laser pulses are 0.6 and 0.8 for the left and right columns, respectively. The ellipticities of the initial transverse momentum at 0 with respect to the ionization time. There is a minimum of the initial transverse momentum for the internal scattered electron calculated by the classical-trajectory model. The ellipticities of the laser pulses are 0.6 and 0.8 for the left and right columns, respectively.

internal scattered electron mainly has two effects. First, it leads to a small spread of the electron wave packet during the propagation. As a result, the internal scattered electrons have substantial contributions to the PMD. Second, the internal scattered electrons only need a small initial transverse momentum to compensate the transverse motion by the elliptical laser field. We show in Fig. 4(c) the absolute value of the initial transverse momentum for the internal scattered electron with respect to the ionization time. There is a minimum of the absolute value of the initial transverse momentum at 0.45T_0, which is earlier than the field maximum. According to the tunneling theory [32], the ionization rate from the rising edge of the laser field is larger than that from the falling edge for the internal scattered electrons. Additionally, we present the final momentum of the internal scattered electron with assuming zero scattering angle. One can see that most electrons are indeed distributed in the first and third quadrants of the momentum plane, which agrees qualitatively with the TDSE results in Figs. 5(c) and 5(d). Thus the internal scattered electrons also have substantial contributions to the PMDs for molecular ionization at the EI region for large ellipticities.

It has been shown recently that the lateral momentum distribution along the laser propagation direction can be dramatically modified by the molecular potential in the EI region [23]. This might also result from the effect of the internal scattered electrons. When the electron wave packet is released from the up-field core and then scattered by the down-field core, the lateral momentum is greatly changed due to the small impact parameter. From Figs. 2 and 3, we can find that the temporal information of the ionization dynamics of H_2^+ has been imprinted in the asymmetry of the PMDs. Comparing the asymmetry of the PMDs of H_2^+ with that of the atom, one can approximately obtain the instant when the electron wave packet is released from the molecule within a half cycle of the laser field. For the internuclear distance of 4 a.u., the electron wave packet is mainly released from the up-field core after the field maximum. As a result, the main emission angle is shifted to the major axis as compared with the atoms. For the internuclear distance of 8 a.u. (EI region), the electron wave packet is mainly released from the up-field core before the field maximum, leading to the opposite asymmetry of the PMD as compared with the atom. This is consistent with the experimental observation in Ref. [25]. Therefore, the asymmetry of the PMD can be used to identify the instant when the electron is released from the molecule in the elliptically polarized laser pulses.

IV. CONCLUSION

We have studied the PMDs of the H_2^+ molecule and a model atom in elliptically polarized laser fields by numerically solving the TDSE. Depending on the internuclear distance of H_2^+, the PMDs reveal different asymmetries with respect to the major and minor axes of the laser ellipse. At an internuclear distance corresponding to the EI region, the asymmetry of the PMD is opposite to that of the model atom. We show that the internal scattered electrons have substantial contributions to the PMDs at the EI region of molecules. By tracing the electron density and using a classical-trajectory model, we find that those internal scattered electrons are mainly released at an earlier ionization moment with respect to the field maximum within each half cycle. Thus the asymmetry of the PMD for molecule at the EI region is opposite to that of the model atom. We further show that the asymmetry pattern guided by the white solid line, which comes from the two-center interference [33]. Those two-center interference fringes are not perpendicular to the molecular axis due to the effect of the molecular potential on the electron dynamics [33]. Besides those two-center interference fringes, we find that most electrons are distributed in the first and third quadrants of the momentum plane, which differs from the PMDs of the atom [Figs. 5(a) and 5(b)]. In Figs. 5(e) and 5(f), we further show the final momentum of the internal scattered electron calculated by the classical-trajectory model with assuming zero scattering angle. One can see that most electrons are indeed distributed in the first and third quadrants of the momentum plane, which agrees qualitatively with the TDSE results in Figs. 5(c) and 5(d).
of the PMD in the elliptical laser fields has recorded fruitful information about the electron dynamics, which could be used to identify the instant when the electron is released from the molecule within a half cycle of the laser field. Our paper provides a deep understanding on the tunneling ionization of molecules in strong laser fields.

ACKNOWLEDGMENTS

This work was supported by the National Natural Science Foundation of China (Grants No. 11627809, No. 11674116, No. 11722432, and No. 61475055) and the Program for HUST Academic Frontier Youth Team.