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Attosecond electro-nuclear dynamics of H₂ double ionization

Sébastien Saugout,^{1,2} Christian Cornaggia,² Annick Suzor-Weiner,¹ and Eric Charron¹

¹*Laboratoire de Photophysique Moléculaire du CNRS,
Université Paris-Sud, Bâtiment 210, F-91405 Orsay, France.*

²*Service des Photons, Atomes et Molécules, Direction des Sciences de la Matière,
CEA Saclay, Bâtiment 522, F-91191 Gif-sur-Yvette, France.*

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The ultrafast electronic and nuclear dynamics of H₂ laser-induced double ionization is studied using a time-dependent wave packet approach that goes beyond the fixed nuclei approximation. The different double ionization pathways are analyzed by following the evolution of the total wave function during and after the pulse. We show that the rescattering of the first ionized electron produces a coherent superposition of excited molecular states which presents a pronounced transient ionic character. This attosecond excitation is followed by field-induced double ionization and by the formation of short-lived autoionizing states which decay via double ionization. These two different double ionization mechanisms may be identified by their signature imprinted in the kinetic-energy distribution of the ejected protons.

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Laser-induced non-sequential double ionization of atoms has been extensively studied since its discovery in the early 1980s [1]. This strong field effect as well as high-order harmonic generation in rare gases [2] gave birth to the so-called rescattering model which gives a unified picture of the atomic response [3]. Moreover, this model constitutes the natural framework for the production of attosecond pulses and for the control of excitation and ionization in atoms and molecules [4]. On the molecular side, new effects in strong laser fields involve the coupling of the electronic and nuclear motions such as bond softening [5] or charge resonance enhanced ionization [6]. Recently, this coupling was shown to play a determinant role in high-order harmonic generation [7, 8, 9]. In particular, Lein predicted that more intense harmonics are generated in heavier molecular isotopes due to the dependence of the harmonics on the vibrational autocorrelation function [7]. This effect has been demonstrated experimentally using H₂, D₂, CH₄ and CD₄ [8]. In addition, molecular vibrations were recently detected with a mÅ sensitivity using high-order harmonic generation in SF₆ [9]. Finally, following a recent theoretical proposal [10], ground state vibrational wave packets of D₂ were mapped with a sensitivity better than 20 mÅ using selective depletion via tunnel ionization [11].

These recent advances show that nuclear motions are fast enough to exercise a decisive influence on the overall electronic response of the molecule in strong fields although the vibrational time scale only belongs to the femtosecond domain. Moreover, the hydrogen molecule exhibits the shortest vibrational periods: 7.5 fs and 14.3 fs for H₂ and H₂⁺. From the theoretical point of view it is therefore highly desirable to develop a unified and realistic theoretical framework where the nuclear dynamics is treated at the same quantum level as the electronic excitation. By treating the internuclear distance R as a

full quantum variable, we show that impulsive vibrational excitation also takes place in H₂. In addition, the rescattering dynamics is basically a two-electron problem which demands a two-active-electron approach. This approach allows for the description of the interplay between electronic rescattering and nuclear vibrations. In our model, we also propose an accurate representation of the field-free molecular electronic states. Using a single-optical-cycle pulse, we show that the electron rescattering leads *during* the pulse to a field-induced double ionization and *after* the end of the pulse to auto-double ionization.

Using the splitting technique of the short-time propagator, we solve the time-dependent Schrödinger equation for H₂ in the linearly polarized classical electric field $\mathbf{E}(t) = E_0 f(t) \cos(\omega t + \varphi) \hat{e}$, where ω denotes the angular frequency, E_0 the field amplitude and φ the carrier-envelope offset phase. The pulse shape is given by the Gaussian-like expression $f(t) = \sin^2(\pi t/\tau)$, with total pulse duration τ . The frequency ω corresponds to a central wavelength of 800 nm. The internuclear coordinate \mathbf{R} is constrained along the field polarization vector \hat{e} and the two electrons, of coordinates $\mathbf{r}_i = z_i \hat{e}$, are assumed to oscillate along the same axis. This one-dimensional approach was shown recently to represent faithfully most of the electronic dynamics [12]. The wave function $\Psi(R, z_1, z_2, t)$ of H₂ is taken initially as the ground state of the soft-Coulomb potential

$$V_c(R, z_1, z_2) = 1/R + [(z_2 - z_1)^2 + \alpha^2(R)]^{-\frac{1}{2}} - \sum_{i=1,2}^{s=\pm 1} [(z_i + sR/2)^2 + \beta^2(R)]^{-\frac{1}{2}} \quad (1)$$

which takes into account the electrostatic repulsion and attraction between all particles. In order to mimic the dynamics of the real H₂ molecule, we have introduced two softening parameters, $\alpha(R)$ and $\beta(R)$, which vary slowly

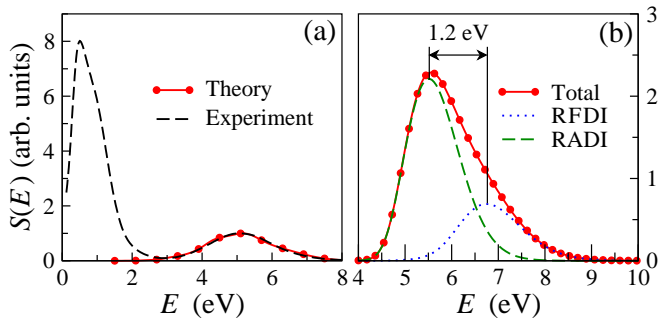


FIG. 1: (Color online) (a) Theoretical (red circles) and experimental (black dashed line) proton kinetic energy spectra for a peak intensity of 4.5×10^{14} W/cm². The measured pulse duration (FWHM) is 10 ± 2 fs and the calculation is for 11.7 fs. (b) Theoretical spectrum (red circles) for a peak intensity of 5×10^{14} W/cm² and a total pulse duration of 2.67 fs. The blue dotted and green dashed lines correspond to recollision-induced field-assisted double ionization (RFDI) and to recollision-induced auto-double ionization (RADI).

with the internuclear distance. These parameters have been adjusted at each internuclear distance to reproduce accurately the potential curves of the ground electronic states of H₂ and H₂⁺. All spectroscopic constants of our model H₂ and H₂⁺ molecules therefore match very closely the tabulated ones. This approach is an extension of the models developed in [13].

During the pulse, the Hamiltonian includes the radiative coupling $(z_1 + z_2)E(t)$ in the dipole length representation. The outgoing ionization flux is summed to extract the single and double ionization probabilities. For this purpose, we partition the plane of the electronic coordinates (z_1, z_2) in three regions Γ_0 , Γ_1 and Γ_2 corresponding respectively to H₂, H₂⁺, and H₂²⁺. Double ionization occurs in the asymptotic region $\Gamma_2 \equiv \{|z_1|, |z_2| > z_A\}$ with $z_A = 20$ a.u. $\gg R$. The single ionization region Γ_1 is defined as $\{|z_1| > z_A \text{ xor } |z_2| > z_A\}$, and the wave function associated with the neutral H₂ molecule is found in the region $\Gamma_0 \equiv \{|z_1|, |z_2| \leq z_A\}$. The total wave function $\Psi(R, z_1, z_2, t)$ is propagated on a spatial grid of $(512)^3$ points with $z_{\max} = 100$ a.u., $R_{\max} = 10$ a.u. and a time step $\delta t = 0.04$ a.u. A simple mapping relates the proton kinetic energy distribution $S(E, t)$ to the probability distribution $P_2(R, t) = \iint_{\Gamma_2} |\Psi(R, z_1, z_2, t)|^2 dz_1 dz_2$ using $E = 1/(2R)$ and the requirement of particle conservation $P_2(R, t) dR = S(E, t) dE$. This energy distribution is accumulated over the entire time propagation to obtain the kinetic energy release spectrum $S(E)$ which is measured in experiments. Fig. 1(a) shows that a good agreement is obtained for the Coulomb peak in the energy range $3 \text{ eV} \leq E \leq 8 \text{ eV}$ between this model and the experiment. Experimental details are given elsewhere [14]. The low energy peak ($E \leq 3 \text{ eV}$), which results from the photodissociation of H₂⁺ in H + H⁺, is of course not reproduced by this Coulomb mapping.

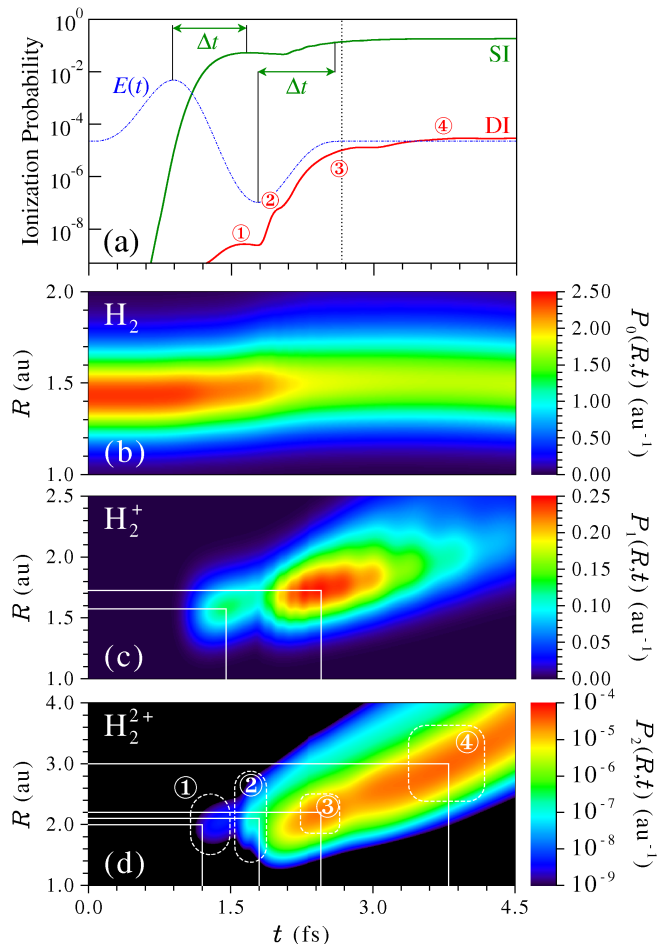


FIG. 2: (Color online) (a) Single (SI) and double (DI) ionization probabilities as a function of time for the total pulse duration 2.67 fs and the peak intensity 5×10^{14} W/cm². The electric field $E(t)$ is superimposed on this graph. The vertical line marks the end of the pulse. (b), (c) and (d) Nuclear probability distributions $P_k(R, t) = \iint_{\Gamma_k} |\Psi(R, z_1, z_2, t)|^2 dz_1 dz_2$ as a function of internuclear distance and time in the three regions Γ_k corresponding to H₂ (b), H₂⁺ (c) and H₂²⁺ (d).

A simplified analysis of the various mechanisms leading to double ionization can be performed theoretically using the shortest pulse containing a complete oscillation of the field, with $\tau = 2\pi/\omega = 2.67$ fs and $\varphi = -\pi/2$. The associated electric field $E(t)$, which is shown Fig. 2(a) as a thin dotted curve, presents two symmetric maxima pointing in opposite directions. The time variations of the single (SI) and double (DI) ionization probabilities are also shown Fig. 2(a) for a peak intensity of 5×10^{14} W/cm². This figure shows that single and double ionization take place by successive bursts. The probability of single ionization is simply rising just after each maximum of the field, and the ionized electron reaches the asymptotic region in $\Delta t \sim 700$ as. The evolution of the double ionization probability is more complex. Indeed, double ionization takes place in four successive bursts which differ consid-

erably by their magnitude and time scale. Since the first appearance of double ionization (label ①, Fig. 2(a)) follows the first maximum of the field by about 700 as, it is clearly due to laser-induced direct double ionization. The second burst (②) just follows the second maximum of the field and thus probably arises from single ionization of the H_2^+ molecular ions already produced. Finally, we will show in the following that the two last and dominant pathways to double ionization (③,④) are induced by recollision and can be attributed to laser-assisted and to autoionization processes.

Fig. 2(b), which represents the evolution of the wave packet in the Γ_0 region, shows that, even for a pulse of total duration $\tau = 2.67$ fs, a significant nuclear motion takes place in H_2 . At about $t = 1.2$ fs, the neutral molecule stretches from the equilibrium internuclear distance $R_e = 1.4$ a.u. to $R \sim 1.5$ a.u. This vibrational wave packet, coherent superposition of a few vibrational levels, arises from the *lochfraß* effect which was predicted and monitored very recently in D_2 [10, 11].

The nuclear dynamics taking place during single and double ionization is even more significant. Figures 2(c) and 2(d) show that the chemical bond stretches up to $R = 1.7$ a.u. when the molecule is singly ionized and up to $R = 3.0$ a.u. during the double ionization process. The first three bursts of double ionization are identified in Fig. 2(d) by the labels ①, ② and ③. These three processes take place at similar internuclear distances, $2.0 \text{ a.u.} \leq R \leq 2.2 \text{ a.u.}$, close to the minimum of the potential associated to the ground electronic state of H_2^+ ($R_e^+ = 2.0$ a.u.). The last mechanism (label ④) is taking place much later, $3.0 \text{ fs} < t < 4.5 \text{ fs}$, after the end of the pulse, and around $R = 3.0$ a.u. This distance is far beyond the equilibrium distances of both H_2 and H_2^+ . In this reaction path, a dissociation process is therefore already taking place during the second stage of ionization, before Coulomb explosion.

A clear identification of the four mechanisms giving rise to Coulomb explosion of H_2 is obtained by analyzing the electronic dynamics shown in Fig. 3 at specific times and internuclear distances. The times t and distances R we have chosen correspond to the four labels given in Fig. 2(d). In agreement with the previous discussion, we observe that laser-induced *direct* double ionization takes place around $t = 1.2$ fs (white arrows in Fig. 3(a)). Indeed, the two electrons escape *simultaneously* from the nuclear attraction in the same direction (z_1 and $z_2 < 0$), opposite to the direction of the electric field. In addition, the electronic repulsion (second term in the *r.h.s.* of Eq. 1) washes out the electronic density around $z_1 = z_2$. The curvature of the two pathways seen in this figure indicates that an energy exchange is taking place, inducing an acceleration of one of the electrons to the detriment of the speed of the other. The electronic probability distribution shown in Fig. 3(b) also confirms that *sequential* double ionization occurs around $t = 1.8$ fs. In this case,

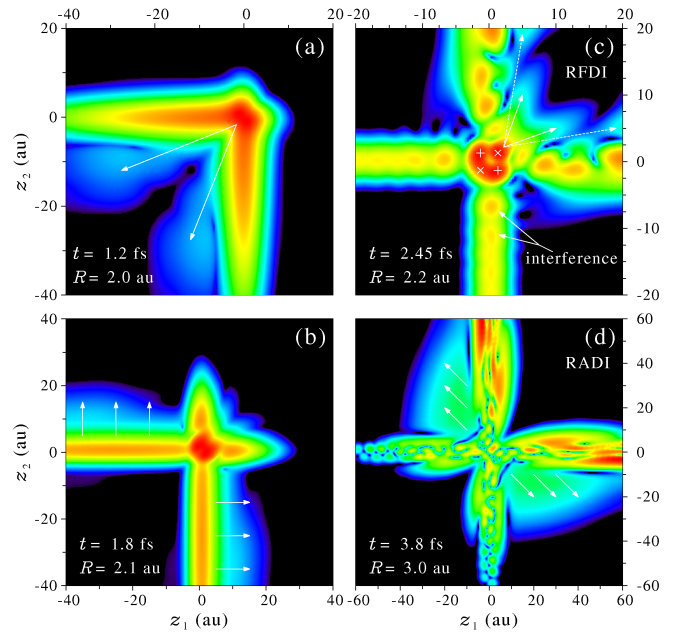


FIG. 3: (Color online) Electronic probability distribution $|\Psi(R, z_1, z_2, t)|^2$ (log. scale) at different times t and internuclear distances R . The laser parameters are the same as in Fig. 2. The white arrows represent the flux of photoelectrons.

the first electron has already left the molecular ion in the direction $z_i < 0$, when the second electron is ejected. The driving electric field has then reversed sign, and this second electron is leaving the nuclei in the opposite direction: $z_j > 0$.

The nature of the two dominant pathways to double ionization is finally enlightened by the snapshots of the electronic density given in the last two panels of Fig. 3. When the electric field changes sign ($t > 1.33$ fs), the part of the electronic wave function which is located in the single ionization region Γ_1 ($z_i < -z_A$ and $|z_j| \leq z_A$), and which therefore represents an ionized electron far from the bound electron of H_2^+ , starts to be driven back towards the nuclei. This recollision process induces an interference between the incoming and outgoing electron waves. This interference is easily seen in the oscillatory behavior of the electronic density in Fig. 3(c). The two peaks identified in this figure, which are separated by $\Delta z \sim 5$ a.u., will recollide successively with the electron that remained near the nuclei in a time interval of about 100 as. Because of this interference pattern, the recollision takes place as a very fast succession of periodic electron impacts. Each impact induces a very specific electronic excitation in the central region where $|z_1|$ and $|z_2| \leq R$. Indeed, in the vicinity of the two nuclei the electronic wave function does not show anymore the usual double peak structure of the ground electronic state of H_2 . This structure is now replaced by a dynamical correlated motion of the two electrons around the four crosses shown in this figure. The two ‘+’ signs correspond to

the two attractive combinations where each electron is located on a different nucleus. The two ‘×’ signs correspond to the two ionic combinations H^+H^- and H^-H^+ . Around the internuclear distance $R = 2.2$ a.u., the energy gap between the ionic state and the ground electronic state of H_2 can be estimated from the energy difference between the $B^1\Sigma_u^+$ and the $X^1\Sigma_g^+$ potential curves of H_2 as $\Delta \sim 0.37$ a.u. Considering the classical ponderomotive energy $U_p = E_{\max}^2/4\omega^2 \sim 0.46$ a.u. for the present maximum value of the field $E_{\max} \sim 0.08$ a.u., the H^+H^- spectral range is clearly accessible following the recollision process. In addition, recent studies [15] have shown that one of these ionic states is stabilized by strong fields, such that the ionic and the ground electronic states of H_2 may experience an avoided crossing in an adiabatic picture. As a consequence, the ionic configuration H^+H^- becomes the instantaneous ground state of the system for large field amplitudes. We find here that this effect, which was first demonstrated in a two-electron model of H_2 with frozen nuclei [15], survives in a more realistic approach which takes into account the dissociative dynamics of the molecule. For large field values, and because of the recollision, the two electrons are thus forced to localize on the same nucleus. Every 100 as, this ionic state is transiently populated by recollision, and in this configuration the two electrons, already on the right hand side of the molecule, can easily escape by tunnel ionization from the nuclear attraction. We therefore observe a very fast succession of bursts of recollision-induced field-assisted double ionization (RFDI, see for instance the four arrows of double ionization in Fig. 3(c)).

As mentioned previously, the last efficient pathway to double ionization takes place after the end of the pulse, and therefore results from an autoionization process of highly excited electronic states. This interpretation is confirmed by the snapshot of the electronic wave packet shown in Fig. 3(d). For $t > 3.0$ fs, the two electrons escape in opposite directions since the ionization is now mainly driven by the electron repulsion. This ionization pathway, as well as the RFDI pathway of Fig. 3(c), are suppressed when we introduce absorbing boundaries for the rescattering flux. The origin of both ionization channels is therefore to be found in the electron recollision.

Finally Fig. 1(b) reports the contribution of these last two ionization channels to the proton kinetic energy spectrum. Since autoionization occurs at larger internuclear distances, the corresponding proton spectrum (dashed line) appears at lower energies than the RFDI proton peak (dotted line). The total spectrum, shown as a solid line with red circles, includes all the double ionization pathways reported in Fig. 2(a). However, the direct and sequential contributions ① and ② remain very weak in comparison with the RFDI and the autoionization mechanisms. The asymmetry of the fragment kinetic energy release spectrum therefore reveals the presence of these

last two pathways, induced by the rescattering process.

To conclude, we have presented a theoretical model for the study of the electronic and nuclear dynamics of diatomic molecules in intense laser fields. This model can be adapted to a particular molecule by a simple adjustment of two softening functions $\alpha(R)$ and $\beta(R)$. We have applied it to the simplest molecular target H_2 . We have shown that at moderate intensities molecular double ionization is mainly driven by the rescattering process. This process induces a specific transient ionic excitation of the molecule which leads to Recollision-induced Field-assisted Double Ionization (RFDI) and to Recollision-induced Auto-Double Ionization (RADI). These two double ionization mechanisms initiate the Coulomb explosion of the molecule at different internuclear distances, an effect which can be observed in the fragment kinetic energy release spectrum. These mechanisms could also be investigated in the time domain using attosecond EUV pulses. Indeed, a time-delayed attosecond pulse may induce an efficient single ionization of H_2 in the energy range $\hbar\omega \sim 20$ eV, thus washing out the electron correlation in a few tens of attoseconds. Controlling the time-delay between the IR femtosecond pump and the EUV attosecond probe should therefore make it possible to reveal experimentally these two mechanisms by removing them one after the other.

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