Light-induced multiple electronic-state coupling of $O_2^+$ in intense laser fields

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The dissociation processes, $O_2^+\rightarrow O^+ + O$, in intense laser fields (100 fs, $\sim 3\times10^{14}$ W/cm$^2$) are studied at the laser wavelength of 795 and 398 nm on the basis of the momentum imaging maps of the $O^+$ fragment ions. The covariance map measurements are performed with high momentum resolution to assign securely the dissociation pathways. From the anisotropic momentum distribution of $O^+$ with respect to the laser polarization direction, the electronic states of $O_2^+$ are found to be coupled exclusively through parallel transitions in the course of the dissociation process. The released kinetic energy of $O^+$ for both 795 and 398 nm are interpreted consistently by a sequential coupling model of light-dressed potential curves of $O_2^+$ in the quartet II state manifold.


I. INTRODUCTION

When the magnitude of a light–molecule interaction exceeds a perturbative regime, interatomic potentials of a molecule are substantially deformed by the light fields. The resultant light-deformed potentials lead to a new class of molecular phenomena such as bond-softerning, bondbreaking, and above-threshold dissociation (ATD). 1 For a one-electron system, $H_2^+$, these phenomena have been interpreted well in terms of the light-dressed picture consisting of the two lowest electronic states, $1\sigma_g$ and $2\pi_u$, both of which are well separated energetically from higher lying electronic states. 1,2 It has been revealed through theoretical calculations 3,4 that the strong coupling between these electronic states is crucial for ionization of $H_2^+$ in intense laser fields, which occurs most efficiently at a critical internuclear distance $R_c \sim 10$ a.u. 3

As for multielectron molecules, the main concern in previous literature has been the multiple-ionization process followed by a bond fission process called the Coulomb explosion occurring at higher field intensities (1–10 PW/cm$^2$; 1 PW = $10^{15}$ W). 5,6 The observed enhancement of the ionization rates at a critical distance has been successfully explained by a classical field ionization model, where the role of electronic states is not taken into account explicitly. On the other hand, recent studies on triatomic molecules 7–12 have revealed that they undergo ultrafast structural deformation in such a high laser-field intensity and that this could be interpreted by the formation of the light-deformed potentials prior to the Coulomb explosion. 9–12 However, such light-induced potential deformation in a multielectron molecule could not be explained by a simple picture in which only two electronic states are coupled with each other as in the case of $H_2^+$, because a multiple coupling would be induced among a number of electronic states with high density.

In the present study, we report definitive evidence of such a multiple coupling among more than two electronic states of $O_2^+$ in intense laser fields ($\sim 0.3$ PW/cm$^2$), through the measurements of the momentum vector distribution of the $O^+$ fragment ions produced from the bond-breaking process of $O_2^+\rightarrow O^+ + O$.

II. EXPERIMENT

The amplified ultrashort laser pulses (795 nm, 100 fs, $\sim 1$ mJ) generated by a regenerative amplifier system (BMI alpha 10) were focused by a quartz lens ($f = 150$ mm) to produce intense laser fields ($\sim 0.3$ PW/cm$^2$). The second harmonics (398 nm) was generated by a BBO crystal plate of 1 mm thickness. The mass-resolved momentum imaging (MRMI) maps of $O^+$ produced from $O_2$ were constructed from the time-of-flight (TOF) mass spectra recorded by an oscilloscope (LeCroy 9370) at 18 different angles of the laser polarization vector with respect to the TOF detection axis in the range between $0^\circ$ and $180^\circ$. The momentum $p_\parallel$ of the ejected fragments along the TOF axis is derived by using a formula, $p_\parallel = qF\Delta T$, where $\Delta T$ is half of the difference in the flight time between the fragments ejected in forward and backward directions, $q$ is the charge of the fragment ion, and $F$ is the static electric field applied for the ion extraction.

A small aperture (10 mm φ) was placed in front of the microchannel plate (MCP) ion detector to achieve high momentum resolution. The pressure in the vacuum chamber was set in the range between $6\times10^{-8}$ and $2\times10^{-7}$ Torr depending on the laser-field intensity to avoid a space charge effect. The energies of the respective laser pulses were monitored by a single channel analyzer, which supplies trigger signals to the oscilloscope. The laser spot size at the focal spot was measured by projecting its image onto a charge coupled device camera through an objective lens ($10\times$) to be $6.4 \times 10^5$ and $1.3 \times 10^5$ μm$^2$ for 795 and 398 nm, respectively.

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In order to discriminate the (1,0) dissociation process $O_2^+\rightarrow O^+ + O$ from the (1,1) Coulomb explosion process $O_2^+\rightarrow O^+ + O^+$, both of which could appear in the MRMI map of $O^+$, the correlation between the $O^+$ fragments was measured by the covariance map technique.$^{13}$ The covariance map was measured after accumulation of more than $3\times 10^4$ shots of laser pulses. The pressure in the chamber was reduced as low as $8\times 10^{-8}$ Torr to keep the ion count rate less than 50 ions/shot. The other experimental conditions were the same as those in the MRMI measurements. Under the present experimental conditions, false covariance signals appeared between $O^+$ ions and $O_2^{2+}$ ions whose magnitude is substantially larger than the $O^+$ signal (see Fig. 1). In order to reduce such an artifact, the raw signal intensities of $O_2^{2+}$ were multiplied by a factor of 0.2 in the construction of the covariance map.

**III. RESULTS AND DISCUSSION**

**A. Assignment of the dissociation pathways**

The observed TOF mass spectra and the covariance map of $O^+$ recorded at $\lambda\sim 795$ nm with the laser polarization vector parallel to the TOF axis are shown in Fig. 1(a). The sharp peak observed at the center of the TOF mass spectra is attributed to $O_2^{2+}$ having the same mass-to-charge ratio as $O^+$. A pair of peaks labeled with a corresponds to the $O^+$ ions ejected in the forward and backward directions with respect to the MCP ion detector. The positive covariance signals between these forward and backward peaks in the covariance map [Fig. 1(a)] indicate that these components are produced in a correlated manner through the (1,1) Coulomb explosion process.$^{14}$ $O_2^{2+}\rightarrow O^+ + O^+$.

Another pair of peaks labeled with b with a small shoulderlike structure on the higher momentum side is seen in the TOF spectrum in Fig. 1(a), exhibiting a marked contrast with $N_2$ (Ref. 15) and NO,$^{16}$ whose TOF spectra showed no clear feature in the smaller momentum region inside the (1,1) pathway. The covariance map of $O^+$ showing no distinct correlation in the small momentum region implies that these slow $O^+$ ions are produced with a neutral O atom as a counterpart fragment through the (1,0) dissociation pathway, $O_2^+\rightarrow O^+ + O$. Existence of such low kinetic energy components of $O^+$ from $O_2$ in intense laser fields was observed previously by Normand et al.$^{14}$ with lower momentum resolution at $\lambda\sim 610$ and 305 nm, and were also assigned to the (1,0) pathway.

The (1,0) component is clearly identified at low laser-field intensities ($\leq 0.1$ PW/cm$^2$) as shown in Fig. 2(a), while the (1,1) component becomes prominent at a higher field intensity ($\geq 0.17$ PW/cm$^2$). The field-intensity dependence shows that the (1,1) Coulomb explosion pathway is composed of two distinct features at $90\times 10^3$ and $105\times 10^3$ amu m/s. By increasing the field intensity further, another pair of peaks appears at the momentum of $140\times 10^3$ amu m/s, originating from the (1,2) Coulomb explosion process, $O_2^{3+}\rightarrow O^+ + O^{2+}$. The lower appearance intensity of the (1,0) component rather than that of the (1,1) component confirms these assignments of the dissociation pathways because a lower charge state of parent molecular ions should be produced with a lower appearance intensity. The present assignment of the (1,0) component is also sup-

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**FIG. 1.** The TOF mass spectrum and the covariance map of $O^+$ (one channel=1 ns) at (a) $\lambda\sim 795$ nm, 0.4 PW/cm$^2$ and (b) $\lambda\sim 398$ nm, 0.7 PW/cm$^2$, obtained for $3\times 10^4$ laser shots. The raw signal intensity corresponding to $O_2^{2+}$ is multiplied by a factor of 0.2 in order to reduce false coincidence signals originating from its strong intensity.

**FIG. 2.** The laser-field dependence of the one-dimensional cut of the MRMI maps of $O^+$ (Fig. 1) along the laser polarization direction for 795 nm (a) and 398 nm (b). The energy scale (in eV) shows the total released kinetic energy.
ported by the measurement of the saturation intensity. From the \(O^+\) yield plotted as a function of the laser field intensity in the 0.1–0.7 PW/cm\(^2\) range, the saturation intensity of the \(O^+\) component is estimated to be 0.2(3) PW/cm\(^2\). The agreement of the saturation intensity with that of \(O_2^+\), 0.2 PW/cm\(^2\), which we estimated from the ion yield obtained by Guo et al.\(^{17}\) at 800 nm, indicates that the slow \(O^+\) ions are produced almost simultaneously with the formation of \(O_2^+\) ion.

The sharp (1,0) peak suggests that only a limited number of the electronic states are involved in the fragmentation process at this field intensity. Since the parent \(O_2^+\) ions are expected to be prepared in its \(b^3\Pi_u\) manifold, an isotropic component with broad momentum distribution in the MRMI map exhibits a large anisotropy along the direction of the laser polarization. As shown in Fig. 3(b), an anisotropic ejection of the \(O^+\) fragment along the laser polarization direction is also identified for the three (1,0) features in the MRMI map observed at 398 nm. In this MRMI map, an isotropic component with broad momentum distribution with a width of \(17 \times 10^3\) amu ms\(^{-1}\) can be seen along the circle with a momentum radius of \(26 \times 10^3\) amu ms\(^{-1}\).

In the ultrafast photodissociation of a diatomic molecule, the fragment anisotropy is determined by the direction of the transition moment \(\mu\) with respect to the molecular axis \(z\).\(^{19}\) When \(\mu/z\) (parallel transition), the angular distribution of the fragment in the space-fixed frame \(P(\theta)\) is expressed as \(P(\theta) \propto \cos^2 \theta\), while in the perpendicular transition \((\mu \perp z)\), \(P(\theta) \propto \sin^2 \theta\), where \(\theta\) is defined as an angle between the laser polarization direction and the direction of the molecular axis.

In the case of a multiphoton transition, the fragment anisotropy can be expressed by a product of these distribution functions \(P(\theta) \propto \cos^{2m} \theta \sin^{2n} \theta\), where \(m\) and \(n\) represent, re-

| TABLE I. The total kinetic energies of \(O^+\) released through \(O_2^+\rightarrow O^+ + O\) in intense laser fields \((-0.1\) PW/cm\(^2\)). The calculated values are derived from the interaction sequence within the quartet-\(\Pi\) manifold. |
|-----------------|-----------------|-----------------|-----------------|-----------------|
|                | 795 nm\(^a\)    | 398 nm\(^b\)    | 610 nm\(^c\)    | 305 nm\(^c\)    |
| 1.1(1)   | 0.8(2)    | 0.5(1)   | 0.6(2)    | 0.72(10)  | 0.7(2)    | 0.52(6)   | —         |
| 2.0(2)   | —         | 1.8(2)   | 1.7(2)    | 1.36(20)  | 2.1(2)    | 1.04(10)  | —         |
| ...      | ...       | 3.7(2)   | 3.6(2)    | 2.60(30)  | —         | 1.94(20)  | 1.5(2)    |
|^aPresent work. The laser pulse duration is \(\tau \approx 100\) fs. |
|^bReference 14 (\(\tau \approx 2\) ps). |
|^cReference 14 (\(\tau \approx 1.4\) ps). |
|^dDominant at a weak field intensity (0.05 PW/cm\(^2\)). |

B. Anisotropic fragment distribution

The observed MRMI map for \(O^+\) recorded at \(\lambda \approx 795\) nm is shown in Fig. 3(a), together with the assignments of the dissociation pathways established above. The (1,0) pathway in the MRMI map exhibits a large anisotropy along the direction of the laser polarization. As shown in Fig. 3(b), an anisotropic ejection of the \(O^+\) fragment along the laser polarization direction is also identified for the three (1,0) features in the MRMI map observed at 398 nm. In this MRMI map, an isotropic component with broad momentum distribution with a width of \(17 \times 10^3\) amu ms\(^{-1}\) can be seen along the circle with a momentum radius of \(26 \times 10^3\) amu ms\(^{-1}\).

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FIG. 3. The MRMI maps of \(O^+\) produced from \(O_2\) in intense laser fields: (a) \(\lambda \approx 795\) nm, \(0.2\) PW/cm\(^2\), (b) \(\lambda \approx 398\) nm, \(0.6\) PW/cm\(^2\). The laser polarization vector \(e\) is indicated with an arrow.
respectively, the number of parallel transitions and that of perpendicular transitions involved in the multiphoton process. Therefore, a nodal line should appear in the MRMI map along the polarization direction when \( n \equiv 1 \), as illustrated in Fig. 4(a), where the simulated MRMI maps\(^{15,16} \) for the three kinds of two-photon dissociation processes, i.e., \( l \perp l ( m = 2, n = 0) \), \( l \perp l ( m = n = 1) \), and \( l \perp l ( m = 0, n = 2) \) are shown. The nodal pattern should remain even when \( \text{O}_2^+ \) is prepared with its molecular axis aligned along the laser polarization direction as shown in Fig. 4(b), in which the alignment distribution is assumed to be expressed as \( P_{\text{align}}(\theta) = \cos^5 \theta \) with respect to the laser polarization vector. The resultant angular distribution is then expressed as \( P(\theta) \cdot P_{\text{align}}(\theta) \). The index \( k \) is chosen to be \( k = 6 \) in the figure in order to approximately describe the fragment anisotropy observed in the MRMI map at \( \lambda \sim 795 \) nm [Fig. 3(a)].

The absence of a nodal line in Fig. 3 means that the electronic states coupled exclusively through parallel transitions dominate the dissociation process. The simulation of the MRMI map, in which an effect of the finite detector size\(^{15,16} \) is taken into account, shows that the observed angular distribution of the \((1,0)\) pathway is well reproduced by the distribution function of \( P(\theta) \approx \cos^5 \theta \) with \( k = 10(4) \) and \( 4(2) \) for \( \lambda \sim 795 \) nm and \( \lambda \sim 398 \) nm, respectively, which are slightly smaller than the \( k \) values for the \((1,1)\) Coulomb explosion pathway obtained from Fig. 3, i.e., \( k = 12(2) \) for \( \lambda \sim 795 \) nm and \( k = 6(2) \) for \( \lambda \sim 398 \) nm. This exclusive choice of parallel transitions may be attributed to the spatial alignment of the molecular axis for the parent ions at the ionization step followed by the strong coupling among the electronic states through charge-resonance type transitions.

The parallel transitions occur among the states with a common electronic symmetry in the limit of the Hund’s coupling case (a). Because \( \text{O}_2^+ \) is expected to be prepared first in either \( X^2\Pi_g \) or \( a^2\Pi_u \), the observed angular distribution suggests that the coupling among the \( \Pi \) electronic states plays a dominant role in the dissociation dynamics of \( \text{O}_2^+ \). This symmetry consideration is important because it reduces significantly the number of electronic states that could be involved in the possible dissociation pathways.

**C. Sequential coupling of the light-dressed states**

The nuclear dynamics of molecules in intense laser fields has often been discussed in terms of light-dressed states \( \{ i, N \} \) that are formed through the coupling between molecular states \( \{ i \} \) and photon-field states \( \{| N \} \). The interaction between molecules and the light fields is expressed as that between the dressed states, leading to potential deformation through avoided crossings. Although this dressed-state picture holds rigorously only when the light field is stationary, it can provide a clear insight into nuclear motions of molecules in the nonstationary strong laser fields.

When \( \text{O}_2^+ \) is prepared in the \( a^2\Pi_u \) state, \( 4\Pi_g \) and \( 4\Pi_u \) electronic states are relevant to the dissociation process, since a transition moment for the transitions among the states with the same spin multiplicity is substantially larger than that for intercombinations. The interatomic potentials of the low-lying \( 4\Pi_g \) and \( 4\Pi_u \) electronic states of \( \text{O}_2^+ \) in the dressed picture are shown in Fig. 5 for the photon energy of 1.56 eV.
The potential curves of the electronic states correlating to the $3P + 4S^o$ and $3D + 4S^o$ asymptotes are taken from a multireference double configuration interaction (CI) calculation by Marian et al.,\textsuperscript{20} where the reliability of the electronic energy was estimated to be better than 0.3 eV. The deviation of the calculated dissociation energy of the $a^4\Pi_u$ state from the corresponding experimental value is only 0.03 eV.\textsuperscript{20} The potential curves of the higher-lying excited states correlating to the $3P + 2D^o$ asymptote are taken from a full valence CI calculation with a minimum-basis set by Beebe et al.\textsuperscript{21}

A large transition moment between a pair of the charge resonance states\textsuperscript{22} could cause a large avoided crossing between the $|a^4\Pi_u,N\rangle$ and $|f^4\Pi_g,N−1\rangle$ potential curves. The dissociation through the (1,0) pathway at $\lambda=795$ nm is thus expected to proceed dominantly from the $v=0$ vibrational levels of the $a^4\Pi_u$ state near the one-photon potential crossing. Since this one-photon crossing is located in energy below the asymptotic level of the $|b^2\Pi_u,N=0\rangle$ state, i.e., $|3P + 4S^o,N=1\rangle$, as seen in Fig. 5, this coupling does not lead directly to the dissociation. The dressed picture in Fig. 5 suggests that additional two dressed states, $|2^4\Pi_u,N=2\rangle$ and $|2^4\Pi_g,N=3\rangle$, located close to the crossing region, can make an interaction sequence $|a^4\Pi_u,N\rangle→|f^4\Pi_g,N−1\rangle→|2^4\Pi_u,N=2\rangle→|2^4\Pi_g,N=3\rangle$, which is expected to occur with a sufficiently large efficiency since it proceeds through a one-photon coupling at each step.

Since the energy of the crossing between $|a^4\Pi_u,N\rangle$ and $|f^4\Pi_g,N−1\rangle$ is located above the dissociation limit of the $|2^4\Pi_u,N=3\rangle$ state, this sequential coupling of the four electronic states leads to the dissociation, which may be referred to as $1+1+1$ ATD. The released kinetic energy for this dissociation pathway, represented by the length of the solid arrow in Fig. 5, is calculated to be 0.8(2) eV from the theoretical energy difference between the $|a^4\Pi_u,N\rangle→|f^4\Pi_g,N−1\rangle$ crossing and the $|1D^+ + 4S^o,N=3\rangle$ asymptote, which is consistent with the observed value of 1.1(1) eV, determined from the momentum distribution of O$^+$ in Fig. 2(a).

At $\lambda=398$ nm, the dressed-state potentials (Fig. 6) are substantially different from those in Fig. 3 due to the doubled photon energy (3.12 eV). As shown in the figure, the coupling among the $a^4\Pi_u$, $f^4\Pi_g$, and $2^4\Pi_u$ states is expected to contribute to the fragmentation as in the case of $\lambda=795$ nm. However, the three $4\Pi_g$ states (3$^2\Pi_g$, 4$^2\Pi_g$, and 5$^2\Pi_g$) correlated to the $3P + 2D^o$ asymptote, denoted hereafter as $4\Pi_g$, may participate in place of the $2^4\Pi_u$ state which is shifted far below in energy from the interaction region. Therefore, the interaction sequence for $\lambda=398$ nm can be expressed as $|a^4\Pi_u,N\rangle→|f^4\Pi_g,N−1\rangle→|2^4\Pi_u,N=2\rangle→|4\Pi_g,N=3\rangle$.

As mentioned in Sec. III B, the MRMI map of O$^+$ at $\lambda=398$ nm [Fig. 3(b)] shows a marked difference from that observed at $\lambda=795$ nm. Apart from the features for the (1,1) Coulomb explosion pathway, three pairs of crescent-like features are observed in the momentum region below 70 $\times 10^3$ amu m/s, with the total release energies of 0.5(1), 1.8(2), and 3.7(2) eV. The appearance of these peaks can be explained by referring to the interaction sequence of the dressed states represented in Fig. 6. At $\lambda=398$ nm, the potential crossing between $|a^4\Pi_u,N\rangle$ and $|f^4\Pi_g,N−1\rangle$ is placed close to the crossing region, denoted hereafter as $|D^+ + 4S^o,N=2\rangle$ and $|3P + 2D^o,N=3\rangle$ states begin to participate in the dissociation process, and lead to the fragmentation into the $|1D^+ + 4S^o,N=2\rangle$ and $|3P^o + 2D^o,N=3\rangle$ asymptotes, respectively. The released kinetic energies for these dissociation pathways represented by the length of the solid arrows in Fig. 6 are calculated to be 0.6(2), 1.7(2), and 3.6(2) eV, which are in good agreement with the corresponding observed energies (see Table I). Therefore, it is reasonable to regard that the three (1,0) peaks $\lambda=398$ nm represent the dissociation into the three different asymptotes.

As seen in Fig. 2(b), the fragments with a larger momentum appear at larger field intensities. This dependence of the peak intensities on the laser-field intensity is consistent with the above interpretation using the coupling sequence of the dressed states at $\lambda=398$ nm. From the intensity variation of these three components in the 0.03–0.7 PW/cm$^2$ range, the saturation intensities were estimated to be 0.10(5), 0.13(5), and 0.18(5) PW/cm$^2$ for the peaks observed at 0.5, 1.8, and 3.7 eV, respectively.

The transition moment $\mu$ of the $f^4\Pi_g→a^4\Pi_u$ charge resonance transition was calculated to be $\mu=0.5$ a.u. at the internuclear distance $R=1.6$ Å, where the $|a^4\Pi_u,N\rangle$ and $|f^4\Pi_g,N=1\rangle$ states cross with each other at $\lambda=398$ nm (see Fig. 5). A slightly larger transition moment, $\mu=0.8$ a.u. was obtained at $R=1.9$ Å for the crossing at $\lambda=795$ nm (Fig. 4).
These values are smaller by a factor of 2–3 than those expected from the formula \( \mu = R/2 \), i.e., \( \mu = 1.5 \) and 1.8 a.u. for \( R = 1.6 \) and 1.9 Å, respectively, because of the configuration mixing of the \( \text{f}^4\Pi_g \) state with other electronic states. Then, the magnitude of the energy gap at the avoided crossing between the charge resonance pair is estimated to be on the order of \( \mu I^{1/2} \sim 1 \) eV at the saturation intensity \( I \sim 0.2 \) PW/cm². Accordingly, an energy gap at a three-photon crossing is expected to be much smaller in the present case. Indeed, the energy gap between the quartet states of \( \text{O}_2^{+} \) is expected to be much smaller in the TOF spectra of \( \text{O}^{+} \) as compared with the observed value by 0.7 eV, the difference is in good agreement with 1.04 eV, one of the remaining two observed values. In order to explain the lowest energy component at 0.52 eV, it may be necessary to take into account additional interactions between the doublet and quartet states in the ionization step that could modulate the initially prepared vibrational distribution in the bound states.

IV. SUMMARY

Based on the MRI measurements of the fragment \( \text{O}^{+} \), the photodissociation process, \( \text{O}_2^{+} \rightarrow \text{O}^{+} + \text{O} \), in intense laser fields \( \sim 1 \) PW/cm², was investigated. The covariance map measurements were performed with high momentum resolution to assign clearly the dissociation pathways. It was found from the anisotropic transitions of \( \text{O}^{+} \) along the parallel and perpendicular vector that the parallel transitions dominate the fragmentation processes, which could be attributed to an anisotropic polarization of the parent ions at the ionization step followed by the strong coupling among the electronic states through charge-resonance type transitions. The total kinetic energy release of \( \text{O}^{+} \) for both 795 and 398 nm was interpreted consistently by a sequential coupling model of light-dressed potential curves of \( \text{O}_2^{+} \) in the quartet II state manifold rather than in the doublet II state manifold. The consistent interpretation of the experimental kinetic energies of \( \text{O}^{+} \) at the two different wavelengths shows that the dissociation dynamics of \( \text{O}_2^{+} \) in intense laser fields \( \sim 1 \) PW/cm² can be interpreted within the framework of the dressed state picture by considering the light-induced multiple coupling among molecular electronic states.

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D. Comparison with previous studies

Normand et al. observed a diffuse peak in the TOF spectra of \( \text{O}^{+} \) at \( \lambda \sim 610 \) nm, 0.05 PW/cm², in which two overlapping components were identified with released kinetic energies of \( \sim 0.7 \) and 1.4 eV. When the present multiple coupling model is adopted, the two interaction sequences, \( \{4 \Pi_u, N \} \rightarrow \{4 \Pi_g, N-1 \} \rightarrow \{4 \Pi_u, N-2 \} \rightarrow \{4 \Pi_u, N-3 \} \) and \( \{4 \Pi_u, N \} \rightarrow \{4 \Pi_g, N-1 \} \rightarrow \{4 \Pi_g, N-2 \} \rightarrow \{4 \Pi_u, N-3 \} \), give the release energies of 0.7(2) and 2.1(2) eV, respectively. Although the latter is slightly larger than the observed value by 0.7 eV, the difference between the observed and calculated peak position is rather small (\( \sim 18 \) ns) in the TOF spectra as compared with the width of the observed diffuse peak (\( \sim 140 \) ns).

The consistent interpretation of the experimental kinetic energies of \( \text{O}^{+} \) at three different wavelengths assures an applicability of the present model of a sequential coupling of the dressed states within the \( 4 \Pi \) manifold and suggests that \( \text{O}_2^{+} \) ion populated in the \( X^2\Pi_g \) state affords only a small contribution to the dissociation process. This may be explained by assuming that the \( X^2\Pi_g \) state is prepared in the low vibrational levels after the ionization step, located far below (\( \sim 6 \) eV) the \( 3P + 3S^0 \) dissociation limit. Since the potential crossing between the charge resonance states, \( X^2\Pi_g, N \) and \( A^2\Pi_u, N-1 \), is located at \( \sim 3 \) and \( \sim 1.5 \) eV above these vibrational levels at \( \sim 795 \) and 398 nm, respectively, the one-photon coupling could not contribute to the dissociation process to a large extent.

On the other hand, the present model within the \( 4 \Pi \) manifold can only account for one of the three components observed at a shorter wavelength at \( \lambda \sim 305 \) nm as shown in Table I. The other two components with lower kinetic energies could be assigned to the dissociation starting from the \( X^2\Pi_g \) state. When the coupling among the doublet II states at \( \lambda \sim 305 \) nm is treated in the same manner as described above, the released energy corresponding to the dissociation into \( \{3P + p\Pi, N-3 \} \) from the crossing between \( X^2\Pi_g, N \) and \( A^2\Pi_u, N-1 \) states is calculated to be \( 1.3(3) \) eV, which is in good agreement with 1.04 eV, one of the remaining two observed values. In order to explain the lowest energy component at 0.52 eV, it may be necessary to account for additional interactions between the doublet and quartet states and/or the resonance effects in the ionization step that could modulate the initially prepared vibrational distribution in the bound states.