I. INTRODUCTION

Oxygen plays a key role in the chemistry of the Earth’s atmosphere. The splitting of the oxygen bond by ultraviolet (UV) solar radiation is the primary step in the formation of ozone (O₃), and is responsible for the existence of the ozone layer. The UV photodissociation of O₂ can be loosely classed in three different wavelength regions: the Herzberg continuum, the Schumann–Runge bands (200–176 nm), and the Schumann–Runge continuum (176–100 nm).

The Herzberg continuum is the region of near-threshold dissociation of O₂ to give O(3P) + O(3P). The electronic configuration of the O atoms (1S²2S²2P⁴) leads to a considerable number of electronic states of the O₂ molecule, and a ground state X 3Σ⁺ that is unusual in that it has two unpaired electrons. In the Herzberg continuum, absorption occurs from the X state to the A 3Σ⁺, A' 3Δᵣ, and c 1Σ⁻ states: these are called the Herzberg I, II, and III transitions respectively. These Herzberg transitions are electric dipole forbidden, and the absorption cross sections (σ) are weak: for example, σ ≈ 10⁻²⁴ cm² at 240 nm. Oscillator strength for the Herzberg transitions is borrowed from electric dipole allowed transitions, mainly through spin–orbit (SO) interactions between ground and excited states. Buijsse et al. have recently published results of a detailed velocity-map imaging study of photodissociation in the Herzberg continuum. The velocity map experiments measured the directions of velocities of the product O atoms with respect to the electric field vector \( \mathbf{e} \) of the linearly polarized photolysis radiation. The angular distribution can be written using the expression:

\[
I(\theta_e) = \frac{1}{4\pi} \{ 1 + \beta P_2(\cos \theta_e) \},
\]

where \( P_2(\cdots) \) is the second order Legendre polynomial, \( \theta_e \) is the polar angle between \( \mathbf{e} \) and \( \mathbf{v} \). The spatial distribution of velocities is parameterized by \( \beta \). The \( \beta \) parameter takes the limits +2 for a parallel (||) transition and −1 for a perpendicular (⊥) transition. At 226 nm, \( \beta = 0.61 \) for O(3P₂), indicating a mixture of 54% (||) with 46% (⊥).

Buijsse et al. have developed a detailed photoabsorption model to directly compare with the experimental results. The Herzberg I transition was found to be the most significant, accounting for 86% of the total cross section at 242 nm, and rising to 94% at 198 nm. The resulting electric dipole transition includes both parallel and perpendicular components:

\[
\begin{align*}
X^{3\Sigma^-}_{g, \pm 1} & \rightarrow B^{3\Sigma^-}_{u, \pm 1} \rightarrow A^{3\Sigma^+}_{u, \pm 1}, \\
X^{3\Sigma^-}_{g, 0+} & \rightarrow 1^{3\Pi^+}_{g, 0+} \rightarrow A^{3\Sigma^+}_{u, \pm 1}, \\
X^{3\Sigma^-}_{g, \pm 1} & \rightarrow 1^{3\Pi^+}_{g, \pm 1} \rightarrow A^{3\Sigma^-}_{u, 0-}.
\end{align*}
\]

Here, we have given the projection \( \Omega = \Lambda + \Sigma \) of total angular momentum on the bond axis as an additional subscript on the term symbol, \( 2\Sigma^± + 1\Lambda_{\Omega, \Lambda} \). The transitions to A' 3Δᵣ and c 1Σ⁻ are purely perpendicular in character. Adiabatically, the A, A', c states correlate to give O atoms with \( j = 2 \), i.e., \( O(3P₂) + O(3P₂) \). Several studies have shown significant populations of \( O(3P_j) \) with \( j = 0, 1, 5, 7 \).
Buijsse et al.\textsuperscript{3} measured the $j=0:1:2$ branching ratio as 1:00:3.33:9.00. Clearly a significant amount of nonadiabatic interaction takes place during the dissociation. There are a total of eight states of ungerade symmetry correlating to the O(3$^3$P$_1$) + O(3$^3$P) dissociation limit, these include three Herzberg states (A, A′, and c) and five others (1$^1$Π$^u$, 1$^3$Π$^u$, 2$^3$Σ$^u$,$^*$$^1$Π$^u$, and 1$^5$Σ$^u$). The ungerade states may couple through long-range SO interaction, and the A$^3$Σ$^u$ and 2$^3$Σ$^u$ states may also interact by radial derivative coupling due to the radial kinetic energy operator. Unfortunately, angular distribution and branching ratio measurements do not reveal clues to the specific states involved in the nonadiabatic dissociation.

Recently, van Vroonhoven and Groenenboom\textsuperscript{8} (VG) calculated \textit{ab initio} potentials for the eight ungerade states, and the SO and radial derivative couplings between them. These potentials were used in semiclassical calculations of the electronic state-resolved branching ratios and $\beta(j)$ parameters as a function of the dissociation energy, and the calculations were compared with the experimental results of Buijsse \textit{et al.}\textsuperscript{9} Significantly, VG also calculated electronic state-resolved alignment moments $\rho^{(k)}_j (k=2,4; j=1,2)$, but at that time no experimental data were available for comparison. The alignment moments are sensitive to nonadiabatic transitions between states, and their experimental determination would provide a useful test of \textit{ab initio} and dynamical calculations.

In this article, we present measurements of electronic orientation and alignment of O(3$^3$P$_1$) atoms following dissociation \textit{via} the Herzberg continuum in the region 218–239 nm. The O-atom polarization is obtained in terms of molecule frame polarization parameters $a^{(k)}_q (p)$ that distinguish between atoms produced by parallel ($\parallel$) or perpendicular ($\perp$) transitions, or coherence due to mixed transitions.\textsuperscript{10} The measured alignment parameters allow a detailed comparison with the recent semiclassical calculations of VG. The photodissociation of O$_2$ with linearly polarized light results in production of O atoms that are oriented in the molecule frame. The orientation results from interference between dissociative states of parallel and perpendicular symmetry. The phase-dependent orientation should prove to be a very sensitive test parameter for the shapes of the dissociative potential energy curves for O$_2$, and the nonadiabatic transitions between states. The paper is arranged as follows: in Sec. II we describe the experiment and extraction of the polarization parameters. The experimental results are presented in Sec. III in comparison with the semiclassical results of VG and our own semiclassical calculations of the interference between excited dissociative states. Conclusions are given in Sec. IV.

II. METHODS

The experimental methods have been described in detail elsewhere,\textsuperscript{11,12} and a brief overview is given here. A 1:9 mixture of oxygen (Matheson, 99.99%) and helium was made and expanded into the source region of a Wiley–Mclaren time-of-flight (TOF) mass spectrometer via a pulsed nozzle (General valve, series 9, 0.6 mm orifice) with a backing pressure of ~400 Torr. The O$_2$ was photolyzed using linearly polarized light in the 218–239 nm wavelength range. The second harmonic (532 nm) of a Nd$^{3+}$:YAG laser (Continuum PL9020) was used to pump a dye laser (Continuum ND6000, rhodamine 590+610 dye) to produce tunable radiation in the region 550–616 nm, which was doubled to 275–308 nm using a KDP crystal. The resulting doubled light was then mixed with the residual fundamental (1064 nm) of the Nd$^{3+}$:YAG laser in a second KDP crystal to produce tunable photolysis radiation from 218 to 239 nm. Measurements of product O(3$^3$P$_j$; j=1,2) atom alignment were made at photolysis wavelengths of 221.667 and 237.049 nm; for brevity, we shall refer to these wavelengths as 222 and 237 nm, respectively. Measurements of O(3$^3$P$_2$) atom orientation were made at several photolysis wavelengths in the range 218.521–238.556 nm. The O(3$^3$P$_j$; j=0,1,2) atom products were ionized by 2+1 REMPI (resonantly enhanced multiphoton ionization) via the 2$p^3$4$p^4$($^3$P$_j$) intermediate state using wavelengths of 226.233, 226.059, and 225.656 nm for j=0, 1, and 2, respectively. Polarization measurements are not possible for O(3$^3$P$_0$) because this state has zero angular momentum. Tuneable probe radiation around 226 nm was produced by frequency doubling the 450 nm output of a second Nd$^{3+}$:YAG pumped dye laser (Spectra Physics DCR-2A, PDL-3, coumarin 450 dye). The probe and photolysis laser beams were counterpropagated and both loosely focused (+50 cm focal length) to overlap at the center of the pulsed expansion in the TOF. The resulting O$^+$ ions were collected with the TOF spectrometer operating under velocity-sensitive conditions.\textsuperscript{12}

The direction of the linear polarization of the photolysis beam was controlled using a double Fresnel rhomb (Optics For Research), and the direction of the probe linear polarization was switched on a shot-by-shot basis using a photoelastic modulator (Hinds PEM 80) allowing difference measurements to be made very accurately. For the orientation measurements, the probe beam was circularly polarized using a Soleil-Babinet compensator (Special Optics), and was switched between left- and right-circular polarization using the PEM. The experimental geometries used were the same as described by Rakitzis \textit{et al.}\textsuperscript{11} The TOF profiles are labeled by their geometry as $I^F_G$, where $F$ indicates the geometry of the photolysis polarization, and $G$ indicates the geometry of the probe polarization. The laboratory Z axis lies along the TOF direction, and the Y axis lies along the photolysis laser propagation direction. Left and right circular polarization is indicated by L and R respectively. For example, for the $I^X_R$ profile, the photolysis laser would be linearly polarized along the X axis, with the counterpropagating probe laser being right circularly polarized. For the alignment experiments, a total set of 4 profiles were recorded. These were obtained in two experiments using the PEM to flip the probe laser polarization between the X and Z axes on a shot-by-shot basis: first with the photolysis polarization along X, and second along Z. This gives two pairs of profiles: \{ $I^X_L$, $I^X_R$ \} and \{ $I^Z_L$, $I^Z_R$ \}. TOF profiles that are sensitive to the O-atom velocity, but approximately independent of product atom alignment are calculated as $I^X_{\text{iso}} = I^X_L + 2I^X_R$ and $I^Z_{\text{iso}} = I^Z_L + 2I^Z_R$. TOF pro-
files that are sensitive to product atom alignment are calculated as $I_{\text{aniso}}^2 = I_{\text{x}}^2 - I_{\text{y}}^2$ and $I_{\text{aniso}}^3 = I_{\text{z}}^3 - I_{\text{x}}^3$. In the orientation experiments, the photolysis radiation was polarized at 45° with respect to the TOF axis, along the ($X-Z$) direction. The probe radiation was switched between left and right circular polarization on a shot-by-shot basis with the PEM to obtain left $I_{\text{L}}^X-Z$ and right $I_{\text{R}}^X-Z$ TOF profiles respectively. Orientation-sensitive and velocity-sensitive composite TOF profiles are defined as $I_{\text{aniso}}^X-Z = I_{\text{L}}^X-Z - I_{\text{R}}^X-Z$ and $I_{\text{iso}}^X-Z = I_{\text{L}}^X-Z + I_{\text{R}}^X-Z$ respectively.\(^{13-16}\)

Following Rakitiz and Zare,\(^10\) we write the relative (2+1) REMPI intensity $I$ as

$$
I[\Theta, \Phi, \theta_e, \beta, a_{q}^{(p)}(p)] = 1 + \beta P_{2}(\cos \theta) + s_{1}(j) \sin \theta \cos \theta, \text{Im}[a_{1}^{(1)}(\|,\perp)] \sqrt{2} \sin \Theta \sin \Phi + s_{2}(j) P_{2}(\cos \Theta) / (1 + \beta) \\
\times \cos^{2} \theta_{e} a_{0}^{(2)}(\|) + (1 - \beta / 2) \sin^{2} \theta_{e} a_{0}^{(2)}(\perp) + s_{2}(j) \sin \theta_{e} \cos \theta_{e} \text{Re}[a_{1}^{(2)}(\|,\perp)] / \sqrt{2}
$$

$$
\times \sin 2 \Theta \cos \Phi + s_{2}(j) (1 - \beta / 2) \sin^{2} \theta_{e} a_{2}^{(2)}(\perp) / \sqrt{2} \sin \Theta \cos 2 \Phi.
$$

\(^{(5)}\)

The molecule frame polarization parameters $a_{q}^{(k)}(p)$ are spherical tensor moments of $j$ in the molecule frame,\(^14\) with the $z$ axis being the recoil velocity vector $v$, assuming that the dissociation occurs in the limit of axial recoil. Equation (5) has been derived by rotating the molecule frame into the laboratory (TOF) frame. The $a_{q}^{(k)}(p)$ are identified by the symmetry of the parent dipole transition moment $(p)$, which is parallel $(\|)$ or perpendicular $(\perp)$ to the diatomic internuclear axis. The symmetry of the transition depends upon the projection of the angular momentum along the O$_2$ bond axis, $\Omega$. For parallel transitions, the change in the projection of angular momentum is $\Delta \Omega = 0$, and for perpendicular transitions $\Delta \Omega = \pm 1$.

The interpretations of the $a_{q}^{(k)}(p)$ are summarized in Table I. The $a_{q}^{(2)}(\|)$ and $a_{q}^{(2)}(\perp)$ represent cylindrically-symmetric alignment of $j$ about $v$ that arise from incoherent parallel and perpendicular excitations respectively.\(^17\) The parameters $\text{Re}[a_{1}^{(2)}(\|,\perp)]$ and $\text{Im}[a_{1}^{(1)}(\|,\perp)]$ arise from the broken cylindrical symmetry of $j$ about $v$ resulting from coherent superposition of parallel and perpendicular excitations. These coherent parameters vanish in the limit of pure parallel or perpendicular transitions. The $a_{q}^{(2)}(\perp)$ results from coherent superposition of perpendicular transitions. The $a_{q}^{(2)}(\perp)$ may occur as a result of simultaneous excitation of $|\Omega| = 1$ states with different energies, or as a result of the excitation of degenerate $\Omega = \pm 1$ components of an $|\Omega| = 1$ transition.\(^17\) It should be noted that relationships between $a_{q}^{(k)}(p)$ and the dynamical functions $f_{k}(q,q')$ of Siebbeles et al.\(^17\) that appeared in Appendix A of Ref. 10 were incorrect. Correct relationships between the molecule frame pa-
I. INTRODUCTION

In our previous works, we have studied the photodissociation of O$_2$ in the ground electronic state at 222 nm and 237 nm. The photodissociation products were measured by time-of-flight mass spectrometry (TOF-MS). In this study, we extend our investigation to the photodissociation of O$_2$ at 226 nm.

II. EXPERIMENTAL 

A. Alignment

For the alignment experiments, the F = X and F = Z profiles are fitted simultaneously, and the sum in Eq. (8) includes $a_{ik}^{(2)}(\parallel)$, $a_{ij}^{(2)}(\perp)$, Re $a_{ij}^{(2)}(\parallel,\perp)$, and Im $a_{ij}^{(2)}(\parallel,\perp)$. For the orientation experiments, F = (X − Z) and Im $a_{ij}^{(1)}(\parallel,\perp)$ is the only polarization parameter fitted by Eq. (8).

III. RESULTS

A. Branching ratios

At the fixed photolysis wavelength 226 nm, we measured the total ion intensity for O($^3P_{1}$) atom REMPI for j = 0:1:2. The ion signals were corrected for variations in probe laser energy. We found the relative branching ratio to be 1.00:2.90:8.65 for j = 0:1:2. Our results are in good agreement with the branching ratios 1.00±0.26:3.33±0.43:9.00±0.70 reported by Buijsse et al. 3 Using branching populations and excitation factors in Tables II and III of VG, 9 we obtain a semicircular estimate at 226 nm of 1.00:4.32:9.73, which agrees well with the experimental results.

B. Alignment

Isotropic and anisotropic TOF profiles $I^F_G$ (F = X, Z) are shown in Figs. 1 (222 nm) and 2 (237 nm) for O($^3P_{2}$) products, and Figs. 3 and 4 for O($^3P_{1}$). The resulting alignment parameters are given in Table II, and plotted in Fig. 5. The incoherent parameter $a_{ij}^{(2)}(\parallel)$ has opposite signs for j = 1 compared with j = 2, and the same is true for $a_{ij}^{(2)}(\perp)$. For a particular j at a given photolysis wavelength, we also find that $a_{ij}^{(2)}(\parallel)$ and $a_{ij}^{(2)}(\perp)$ have opposite signs. The $a_{ij}^{(2)}(\parallel)$ does not appear to change, within uncertainties, from 222 nm compared to 237 nm and the $a_{ij}^{(2)}(\perp)$ appears to decrease at
TABLE II. Experimental beta parameters ($\beta$) and molecule frame alignment parameters $a^{(2)}(p)$ for O(3P) atoms at photolysis wavelengths 222 and 237 nm. The calculated semiclassical $a^{(2)}(p)$ from Table III are also shown here for convenience. The calculated $\text{Re}[a^{(2)}(\langle\|\rangle)]$ and $a^{(2)}(\langle\|\rangle)$ were obtained using methods outlined in Sec. III C.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>222 nm</th>
<th>237 nm</th>
<th>222 nm</th>
<th>237 nm</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\beta$</td>
<td>0.51±0.10</td>
<td>0.25±0.15</td>
<td>0.55±0.10</td>
<td>0.50±0.10</td>
</tr>
<tr>
<td>$a^{(2)}(||)$</td>
<td>+0.158±0.043</td>
<td>+0.123±0.048</td>
<td>-0.090±0.037</td>
<td>-0.125±0.030</td>
</tr>
<tr>
<td>calculated</td>
<td>+0.055</td>
<td>-0.007</td>
<td>-0.060</td>
<td>-0.191</td>
</tr>
<tr>
<td>$a^{(2)}(\langle|\rangle)$</td>
<td>-0.223±0.071</td>
<td>-0.052±0.057</td>
<td>+0.081±0.066</td>
<td>+0.015±0.050</td>
</tr>
<tr>
<td>calculated</td>
<td>-0.131</td>
<td>-0.048</td>
<td>+0.073</td>
<td>+0.080</td>
</tr>
<tr>
<td>$\text{Re}[a^{(2)}(\langle|\rangle)]$</td>
<td>-0.115±0.042</td>
<td>+0.069±0.035</td>
<td>+0.090±0.037</td>
<td>-0.001±0.027</td>
</tr>
<tr>
<td>calculated</td>
<td>+0.019±0.035</td>
<td>+0.061±0.023</td>
<td>-0.094±0.033</td>
<td>-0.009±0.021</td>
</tr>
<tr>
<td>$a^{(2)}(\langle|\rangle)$</td>
<td>+0.019±0.035</td>
<td>+0.061±0.023</td>
<td>-0.094±0.033</td>
<td>-0.009±0.021</td>
</tr>
</tbody>
</table>

The molecule frame alignment tensor moments $\rho^{(2)}$ calculated by VG (Ref. 9) can be related to the molecule frame alignment parameters $a^{(2)}(\|\|)$ and $a^{(2)}(\langle\|\rangle)$ of Rakitzis et al.,

$$a^{(2)}(\|\|) = \left[ \frac{(2j-1)(2j+1)(2j+3)}{5(j+1)} \right]^{1/2} \rho^{(2)}.$$  

(9)

The semiclassical calculations of VG were obtained at energies equivalent to photolysis wavelengths 204, 216, 226, 236, and 240 nm and our experimental results were obtained at 221.667 and 237.049 nm. We have made linear interpolations of the results of VG in order to compare with our experimental data: these were calculated as $\rho^{(2)}(222\text{ nm}) = 0.5667 \times \rho^{(2)}(226\text{ nm}) + 0.4333 \times \rho^{(2)}(216\text{ nm})$, and $\rho^{(2)}(237\text{ nm}) = 0.7377 \times \rho^{(2)}(236\text{ nm}) + 0.2623 \times \rho^{(2)}(240\text{ nm})$. The resulting $\rho^{(2)}$ were converted to $a^{(2)}(\|\|)$, using Eq. (9), and are given in Table III. Also shown in Table III are interpolated semiclassical excitation branching ratios to each of the Herzberg states, and their respective $\beta$ parameters.

From Table III we see that dissociation via $A^3\Sigma^+_u,1$ has both parallel and perpendicular contributions. For example, if $\beta=1.219$ we determine that $(\beta+1)$ of $0.740$ is the parallel branching fraction of the $A^3\Sigma^+_u,1$ state dissociation at 222 nm. The total parallel branching fraction for O$_2$ dissociation at 222 nm is $0.735 \times 0.740 = 0.544$. From Table III, $a^{(2)}(\|\|)$ is $-0.111$ for the $A^3\Sigma^+_u,1$ state at 222 nm. We calculate the total parallel contribution to alignment as $a^{(2)}(\|\|) = -0.111 \times 0.544 = -0.060$. The last two rows in Table III are the weighted sums of the contributions from parallel and perpendicular transitions, and the calculated $a^{(2)}(\|\|)$ results are compared to the experimental results in Table II. Comparing the experimental and calculated $a^{(2)}(\|\|)$ in Table II, we see good quantitative agreement overall, within experimental uncertainties. There appears to be greater discrepancy between experimental and calculated results for $a^{(2)}(\|\|)$, particularly for $j=1$ atoms.

The dissociation of O$_2$ via the Herzberg continuum approaches the threshold for the O(3P)+O(3P) dissociation limit, and we must consider the potential effects of nonaxial recoil. In the $X^1\Sigma^+_g$ electronic state there are only odd rotational $J$ states, with the lowest populated level being $J=1$. The estimated rotational temperature of our expansion is 15 K, and we calculate fractional Boltzmann populations as $0.787, 0.197,$ and 0.016 for $J=1, 3,$ and 5, respectively. The dissociation energy of O$_2$ is $D_0(O_2) = 41268.6\text{ cm}^{-1}$. Assuming dissociation from O$_2(J=1)$ to give two ground state
TABLE III. Herzberg excitation branching ratios $p_{n\Omega}$ and state-specific anisotropy parameters $\beta_{n\Omega}$ for different molecular eigenstates, calculated from Table II of VG (Ref. 9). Semiclassical alignment parameters $a_0^{(2)}(j=1,2)$ were calculated by interpolation from the results of VG, see text for details.

<table>
<thead>
<tr>
<th>State</th>
<th>$p_{n\Omega}$ 222 nm</th>
<th>$p_{n\Omega}$ 237 nm</th>
<th>$\beta_{n\Omega}$ 222 nm</th>
<th>$\beta_{n\Omega}$ 237 nm</th>
<th>$a_0^{(2)}(j=1)$ 222 nm</th>
<th>$a_0^{(2)}(j=1)$ 237 nm</th>
<th>$a_0^{(2)}(j=2)$ 222 nm</th>
<th>$a_0^{(2)}(j=2)$ 237 nm</th>
</tr>
</thead>
<tbody>
<tr>
<td>$A^1\Sigma^+_u,1$</td>
<td>0.735</td>
<td>0.727</td>
<td>1.219</td>
<td>1.253</td>
<td>0.102</td>
<td>0.012</td>
<td>0.111</td>
<td>0.350</td>
</tr>
<tr>
<td>$A^1\Sigma^+_u,0$</td>
<td>0.191</td>
<td>0.181</td>
<td>1</td>
<td>1</td>
<td>1.648</td>
<td>0.154</td>
<td>0.378</td>
<td>0.235</td>
</tr>
<tr>
<td>$e^1\Sigma^+_g^-$</td>
<td>0.021</td>
<td>0.027</td>
<td>1</td>
<td>1</td>
<td>0.174</td>
<td>0.254</td>
<td>0.435</td>
<td>0.271</td>
</tr>
<tr>
<td>$A^1\Delta^+_u,1$</td>
<td>0.021</td>
<td>0.026</td>
<td>1</td>
<td>1</td>
<td>0.174</td>
<td>0.254</td>
<td>0.435</td>
<td>0.271</td>
</tr>
<tr>
<td>$A^1\Delta^+_u,2$</td>
<td>0.031</td>
<td>0.039</td>
<td>1</td>
<td>1</td>
<td>0.174</td>
<td>0.254</td>
<td>0.435</td>
<td>0.271</td>
</tr>
<tr>
<td>$A^1\Delta^+_u,3$</td>
<td>0.001</td>
<td>0.001</td>
<td>1</td>
<td>1</td>
<td>0.174</td>
<td>0.254</td>
<td>0.435</td>
<td>0.271</td>
</tr>
<tr>
<td>total $\parallel$</td>
<td>0.544</td>
<td>0.546</td>
<td>2</td>
<td>2</td>
<td>0.155</td>
<td>0.007</td>
<td>0.060</td>
<td>0.191</td>
</tr>
<tr>
<td>total $\perp$</td>
<td>0.456</td>
<td>0.454</td>
<td>1</td>
<td>1</td>
<td>0.101</td>
<td>0.048</td>
<td>0.073</td>
<td>0.000</td>
</tr>
</tbody>
</table>

O($^1P_2$) atoms, we estimate the final center-of-mass velocity of O atoms to be 1681 m s$^{-1}$ at 222 nm and 832 m s$^{-1}$ at 237 nm. Following Wrede et al., we calculated the total quasiclassical angle of rotation $\gamma$ that accumulates during the dissociation, i.e., $\gamma$ is the angle between the asymptotic velocity vector of the O atoms and the position of the internuclear axis at the time of excitation. For O$_2$ ($J=1$) we find that $\gamma=1.4^\circ$ at 222 nm and $\gamma=2.5^\circ$ at 237 nm, and for O$_2$ ($J=3$) $\gamma=3.5^\circ$ at 222 nm and $\gamma=6.2^\circ$ at 237 nm. The effect of these deviations from axial recoil are small: for $\gamma=6.2^\circ$ $\beta$ would be degraded to only 0.98 of its axial value.

A further complication in our experiment is that we are unable to determine the correlated alignment between $j_A$ and $j_B$ for the two atoms, so there is an implicit averaging over the co-fragment with $j=0,1,2$. The energy separation $\approx 100$ cm$^{-1}$ between $j$ levels of O($^3P$) would make it very challenging to obtain correlated experimental data. It is possible, however, that the effects of the implicit averaging could be assessed in future calculations.

In summary, the semiclassical $a_0^{(2)}$ of VG agree well with our experimental results. We expect that a fully quantum model of the Herzberg excitation and dissociation, which could include calculation of the coherent alignment parameters, would improve the agreement between theory and experiment.

C. Orientation

Typical isotropic and anisotropic TOF profiles $I_{\Omega}^{Z-Z}$ at photolysis wavelength 223.543 nm are shown in Fig. 6. The resulting $\text{Im}[a_1^{(1)}(\|,\perp)]$ obtained from measurements over a range of photolysis wavelengths 218.521–238.556 nm are shown in Fig. 7, and tabulated in Table IV. The $\text{Im}[a_1^{(1)}(\|,\perp)]$ parameter represents orientation along the molecule-frame $y$ axis resulting from interference between states of parallel and perpendicular symmetry. The $\text{Im}[a_1^{(1)}(\|,\perp)]$ contains both amplitude and phase information. For the simple case of interference between two states (one parallel, one perpendicular) we can write

$$\text{Im}[a_1^{(1)}(\|,\perp)] \propto \sin \Delta \phi,$$

where $\sin(\Delta \phi)=\sin(\phi_1 - \phi_2)$ is the sine of the asymptotic phase difference between the radial parts of the dissociative wave functions for the perpendicular and parallel states. The photodissociation of O$_2$ involves more than one pair of interfering electronic states. A fully quantum mechanical simulation of the dissociation is outside the scope of the present study. However, as a base to compare with our experimental results, we have calculated the energy dependence of the semiclassical phase difference using the O$_2$ potential energy curves of VG.

We begin by writing the $\text{Im}[a_1^{(1)}(\|,\perp)]$ in terms of the dynamical functions $f_1(q,q')$ of Siebbeles et al.

$$\text{Im}[a_1^{(1)}(\|,\perp)] = -\frac{6 \text{Im}[f_1(1,0)]}{\sqrt{2}[f_0(0,0) + 2f_0(1,1)]}.$$  

The dynamical functions can be written as

![Figure 6](https://example.com/fig6.png)

**FIG. 6.** Isotropic and anisotropic TOF profiles of O($^3P_2$) products at photolysis wavelength 223.543 nm: (a) $I_{\text{iso}}^{Z-Z}$, (b) $I_{\text{aniso}}^{Z-Z}$. Experimental data are shown as dashed lines, fitted data are solid lines.
The dissociation gives two fragments with angular momenta $j_A$ and $j_B$, having projections $\Omega_A$ and $\Omega_B$ on the molecule frame $z$ axis. The indices $n$ label the excited state with $\Omega = \Omega_A + \Omega_B$. To be brief, we shall refer to the six Herzberg states of Table III as $A_1$, $A_0$, $c_0$, $A'_1$, $A'_2$, and $A'$. As explained by Balint–Kurti et al., summation over $\Omega_B$ is implicitly included in Eq. (12), and $q - q' = \Omega_A - \Omega_A' = \Omega - \Omega'$. The coefficients $\mathcal{T}^{0}_{j_Aj_Bj_A'j_B'}$ are expansion coefficients of the molecular electronic states in the product atomic basis $|j_A\Omega_A,j_B\Omega_B\rangle$, which can be calculated by diagonalizing the long-range quadrupole–quadrupole potential interaction matrix in the atomic basis. $\mathcal{T}^{0}_{j_Aj_Bj_A'j_B'}$ coefficients for O$_2$ have been given in Table I of VG.\(^9\)

The dynamical function $f_1(1,0)$ represents a sum over (||,) pairs of interfering states. No interference occurs between dissociation paths that start with different values of the ground state $\Omega$. Since the only parallel transition involved is from ground state $X_1 \rightarrow A_1$ [see Eq. (2)], the only perpendicular transitions to be considered in $f_1(1,0)$ are $X_1 \rightarrow A_0$, $c_0$, and $A'$.\(^2\)

The coefficients $r_{n\Omega}$ and $\phi_{n\Omega}$ in Eq. (12) are amplitudes and phases of the photofragment T matrix. For their work on HF photodissociation,\(^{21}\) Balint–Kurti et al. calculated the T matrix from time-dependent wave-packet dynamical calculations. In the present case, we approximate by using the semiclassical results of VG to obtain amplitudes as the (positive) root of the branching ratios in Table III. For parallel transitions $r_{n\Omega} = (P_{n\Omega})^{1/2}$, and for perpendicular transitions $r_{n\Omega} = (P_{n\Omega}/2)^{1/2}$. The branching ratios $P_{n\Omega}$ depend on the photolysis wavelength, and are given in Table II of Ref. 9. The parallel component $P_{A1\Omega} = P_{A1}(1 + \beta)/3$ of the branching fraction to $A_1$ has been used. Taking the positive root we have introduced a factor of $\pi$ uncertainty in the phase for each pair of interfering states in $f_1(1,0)$.

The total semiclassical phase $\phi_n$ for state $n = n\Omega$ was found by numerical integration using the adiabatic Hund’s case (c) potential energy curves of VG with the expression\(^{23}\)

$$\phi_n = \frac{\pi}{4} + \int_{R_{n}^{(2)}(\Omega_2)}^{\infty} \left[ k_n(R) - k_n(\infty) \right] dR - k_n(\infty) T_n,$$

where $R$ is the O$_2$ bond length, $T_n$ is the classical inner turning point for state $n$ at a given available energy $E$. The available energy $E = h\nu - D_{0}^{0}(O_2)$, where $h\nu$ is the energy of the photolysis photon. The de Broglie wave number $k_n(R)$ is

$$k_n(R) = \frac{1}{\hbar} \sqrt{2\mu [E - E_n(R)]},$$

where $\mu$ is the reduced mass of O$_2$, and $E_n(R)$ is the potential energy of state $n$. The centrifugal potential due to rotation of the dissociating O$_2$ was found to be insignificant in our calculations and has not been included in Eq. (14). The case (c) potentials were obtained by diagonalization of the matrix elements of the Hamiltonian $\hat{H} = \hat{H}_{\text{Col}} + \hat{H}_{\text{SO}}$ in the basis of the electronic states for $\Omega = 0, 1, 2,$ and 3, respectively, as a function of the internuclear separation, $R$. The eigenvalues of the Coulombic Hamiltonian $\hat{H}_{\text{Col}}$ correspond to the electronic adiabatic Born–Oppenheimer (ABO) states, and the matrix elements of $\hat{H}_{\text{SO}}$ are the spin–orbit couplings between ABO states.\(^{24}\)

Equations (11)–(14) show how $\text{Im}[a_{1}^{(1)||,\perp}]$ is related to the potential energy curves $E_n(R)$ through the phase accumulated during dissociation at a given available energy $E$. When the available energy is much larger than the potential energy difference between parallel and perpendicular curves, the phase difference $\Delta \phi$ does not vary much, and the $\text{Im}[a_{1}^{(1)||,\perp}]$ will change very slowly as a function of the photolysis energy. Nearer to threshold, however, $\Delta \phi$ varies more rapidly with $E$, and measurement of the energy dependence of $\text{Im}[a_{1}^{(1)||,\perp}]$ can be a very sensitive test of the shapes of the potential energy curves.\(^{25}\) For the present case, we calculate

![FIG. 7. Molecule frame orientation parameter Im[a_{1}^{(1)||,\perp}]](nm) vs photolysis wavelength. Circles are experimental points. The solid line is the semiclassical simulation obtained using potential energy curves of VG [see Eq. (15) (Ref. 8)]. The dashed line incorporates a reduction in the term involving the $c_0$ state. See text for details.](Image)
\[
\text{Im}[a_i^{(1)}(\|,\perp)] = f_{A0} r_{A0} r_{A11} \sin(\Delta \phi_{A0}) \\
+ f_{c0} r_{c0} r_{A11} \sin(\Delta \phi_{c0}) \\
+ f_{A12} r_{A12} r_{A11} \sin(\Delta \phi_{A12}),
\]
where \(\sin(\Delta \phi_{A1}) = \sin(\phi_{A1} - \phi_{A1})\) and the constants \(f_{A0} = -1.060, f_{c0} = -1.325,\) and \(f_{A12} = -1.458.\) Recall that we have no knowledge about the sign of each term in Eq. (15). We have therefore chosen the coefficients \(f_{A1}\) to have the same sign, which qualitatively matches the experimental data. The results of the semiclassical calculation of Eq. (15) are shown as the solid line in Fig. 7. Even with uncertainty in the relative signs of the three terms in Eq. (15), the magnitude of the calculated \(\text{Im}[a_i^{(1)}(\|,\perp)]\) is remarkably close to the experimental points. There are pronounced ripples in the orientation: the frequency of the ripples increase at longer wavelength, as the available energy decreases on approach to the threshold. At wavelengths around 225 nm, the magnitude of oscillation in the calculated orientation is larger than experimentally observed. The oscillations in this energy region are mostly due to the interference component involving the \(c0\) state. By fitting the experimental data using a nonlinear least-squares algorithm, we investigated the effect of reducing the contribution from the \(c0\) state. We find that the experimental results are more closely followed if the contribution from the \(c0\) state is reduced by multiplying the second term in Eq. (15) by a constant factor of 0.15. The resulting orientation is shown as the dashed curve in Fig. 7. The reduction factor is equivalent to reducing the branching fraction \(p_{c0}\) in Table III at 222 nm from 0.021 to \(4.5 \times 10^{-4}\). This is a sizable reduction, and it is very doubtful that this is the sole source of discrepancy between the experiment and our semiclassical calculations. Unfortunately, the maximum observed in the calculated results at around 230 nm can only be inferred in our experimental data as it covers a region that was difficult to access with our photolysis nonlinear mixing setup. Despite the simplicity of our semiclassical calculations, the results show reasonable agreement with the experimental results.

**IV. CONCLUSIONS**

We have made a detailed study of the \(3P_J\) atom alignment and orientation following photodissociation of \(O_2\) in the Herzberg continuum in the region 218–239 nm. The polarization parameter formalism used has allowed us to determine alignment contributions from parallel or perpendicular transitions, and alignment resulting from coherence between excited dissociative states. Experimental alignment parameters were found to be in reasonably good agreement with recent semiclassical simulations by van Vroonhoven and Groenenboom (VG). We have found that photolysis with linearly polarized light produces oxygen atoms that are oriented as a result of interference between multiple excited dissociative states of parallel and perpendicular symmetry. The photolysis energy dependence of the orientation can be related to the phase difference between parallel and perpendicular states. Semiclassical calculations of the energy-dependent orientation were made using Hund’s case (c) potential energy curves of VG, and were shown to give good agreement with the experimental results.

The photodissociation of \(O_2\) via the Herzberg continuum reveals a surprisingly rich photochemistry involving motions on—and interactions between—more than one electronic potential energy surface that correlates to the separated open-shell atomic photofragments. This work brings out the inadequacy of the Born–Oppenheimer approximation in describing the dissociation of simple molecules when the dissociating state lies in the midst of other excited states of different symmetry. The Herzberg continuum serves as a fundamental example of nonadiabatic photodissociation. As further theoretical studies are carried out, we hope that our experimental results will provide additional tests for our understanding of this important atmospheric process.

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