Predissociation of $O_2$ in the $B$ state

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Dispersed LIF spectra of $O_2$ in the Schumann–Runge band were measured with a modified tunable ArF laser in a flame. Spin-state selective predissociation of the $B$ state was directly observed in fluorescence excitation spectra, revealing the relative coupling matrix elements of the triplet components to the manifold of repulsive states. Such data determines the symmetries of the important predissociating curves for each observed $B$-state vibrational level and shows that past interpretation of absorption linewidth data is in error. Due to the fast predissociation, quench-free emission spectra arising from laser prepared single rovibronic levels in the $B$ state were observed even in an atmospheric flame. Fluorescence to $X$-state vibrational levels as high as $v' = 35$ was observed and relative emission probabilities were derived.

I. INTRODUCTION

The spectroscopy of $O_2$ in the Schumann–Runge band is one of the most extensively studied topics in diatomic spectroscopy. The importance of $O_2$ in solar absorption, atmospheric photochemistry, and combustion, as well as being a colorful example of molecular structure, has provided the motivation for the extensive effort given to this problem.

The Schumann–Runge band is photochemically very important because of the predissociation of the $B$ state. Absorption linewidth measurements have shown vibrational and rotational state dependent lifetimes for $v' = 0–19$ and $J' = 0–20$, varying from 2 to 25 ps. Such data has been the subject of extensive theoretical effort. Spin–orbit coupling of the $B$ state to a manifold of repulsive states is the dominant predissociation mechanism. The vibrational state dependence for the $J = 0$ states has been fit and interpreted as the result of four repulsive curves, each with different vibrational state dependent predissociation efficiencies. According to this analysis, the $1\Pi_u$ repulsive curve, which is also responsible for the observed spectroscopic perturbations analyzed by Julienne, is dominant for nearly all vibrational levels. The other three ($1\Pi_u$, $3\Pi_u$, and $1\Sigma_u^+$) are thought to play a minor but nonnegligible role and are predicted to produce no spectral perturbations, observable with available resolution. The rotational state dependence has been fit assuming a simple centrifugal distortion model within the spin–orbit dissociation mechanism. No evidence of orbit–rotation interaction is seen in the rotational dependence.

Despite the good theoretical fit to the results of absorption spectra, one must view this apparently well-treated problem with suspicion. The spin–orbit operator represents an interaction of the magnetic dipole moment of the electron spin with a magnetic field due to electronic angular momentum. An interaction between different degrees of freedom of electronic motion may not produce clearly interpretable rotational and vibrational dependent predissociation lifetimes. The clearest effect should appear in the spin-state dependence of the lifetimes. Such experimental results are not yet available for $O_2$ from absorption spectra.

Dispersed LIF spectra of $O_2$ exist in only a few cases, because of the very low fluorescence quantum yield and the lack of intense light sources in the wavelength region of the Schumann–Runge band. This is however a much better way to investigate the state dependent predissociation of $O_2$ in the $B$ state. By dispersing the fluorescence, the spectra can be simplified and as will be shown the triplet can be resolved. In LIF there is no need to make linewidth measurements because the signal intensities are directly proportional to the predissociation lifetimes.

In this experiment strong LIF spectra of $O_2$ have been observed using a tunable ArF laser. The wavelength scanning range of this laser is sufficient to interrogate transitions to $v' = 10, 11, 12, 14, 15, 16$, and 17. For $v' = 11, 14, 15$, and 16, the resolution of the laser was sufficient to resolve the components of the spin multiplets. This provides a very clear picture of the predissociation mechanism in $O_2$.

The high pumping efficiency of the laser has also allowed the observation of dispersed emission spectra of $O_2$ originating from $v' = 10, 11, 14, 15$, and 16 and terminating in states as high as $v'' = 35$. The relative emission probabilities between vibrational states of the $B$ and $X$ curves have been extended nearly to the dissociation limit of the $X$ state.

II. EXPERIMENTAL

Figure 1 illustrates the experimental arrangement. In order to populate the states of interest, $O_2$ was observed near the edges of a flame, produced by a butane soldering torch that was borrowed from the workshop and used without modification. There are no collisional effects in the experiment because the predissociation rate of the $B$ state is much faster than the intermolecular collision rate in the atmospheric flame. This can be seen clearly in the emission spectra shown below (Sec. IV). Laser induced fluorescence was produced by the unfocused output of a modified Lambda-physics EMG150 tunable excimer laser operating with argon fluoride. The laser delivered light pulses $3 \times 1$ cm in size, with 100 mJ of energy and a $1 \text{ cm}^{-1}$ bandwidth. One part in 1000 of the LIF was collected by an $f/5.3$ one-to-one imaging lens and was dispersed through a 0.3 m Macpherson 218 monochromator with a linear dispersion of 2.65 nm/mm and 0.2 $\times$ 10 mm slits. The estimated volume of the flame sampled by the imaging system of the monochromator was...
The narrow band output of the oscillator was measured at a 2.4 m distance, the distance between the laser output and the grating. To include the laser output as a function of wavelength, the spectra were simultaneously recorded by a strip-chart recorder and stored in the memory of an IBM PC. The transmission function of the monochromator was sufficient to remove all scattered laser light away from the narrow band output of the oscillator. We have observed only very weak transitions from accessible upper states at the ArF wavelength.9 Excitation of states above \( v'' = 3 \) goes directly to the B-state continuum and is expected to play no role in this experiment.

Figure 2 shows a typical LIF spectrum of \( \text{O}_2 \) in a flame with a spectral simulation (above). The monochromator was adjusted to observe \( v' = 14 \rightarrow v'' = 21 \) emission at 351.5 nm with 0.5 nm resolution. This excitation spectra results primarily from the \( v' = 14 \rightarrow v'' = 3 \) transition, although \( v' = 15,16 \rightarrow v'' = 3 \) are also present. This signal to noise was obtained with a scan duration of about 10 min. The relative intensities between rotational states could be fit with a temperature of 1750 ± 250 K. This is in agreement with previous temperature measurements of \( \text{OH} \) using a tunable KrF laser.8 Spatially dependent temperature measurements in Ref. 6 also showed that there is only a small temperature gradient in the region of the flame probed by this experiment.

The apparent doublets in the spectrum are due to the partial resolution of the \( \text{O}_2 \) spin triplet in \( v' = 14 \). Each component of the triplet has the same \( N \) quantum number (referring to nuclear rotor angular momentum). The red shifted transition always prepares the B-state \( F_i \) component, scanning range and better locking efficiency. The details and quantitative improvement achieved will be reported in a separate publication.7 The principal modification was to replace the 1.5 mm holes that define the active volume of the oscillator cavity with 1.5 X 5 mm slits and to telescope the resulting rectangular output of the oscillator back to a 1.5 X 1.5 mm spot using two cylindrical lenses. This improved not only the output power of the oscillator but also reduced the shot-to-shot fluctuations that arise from instabilities in the discharge.

The shallow valley which underlies all the spectra shown below is an indication that the laser is operating well. On the edges of the scanning range where the broad band component of the excimer laser is a large part of its total emission, many \( \text{O}_2 \) transitions are weakly excited and observable at the chosen emission wavelength. As the laser becomes well locked in the middle of the scanning range, this source of signal diminishes to about 7% of the resonant signal. A comparison of the background noise level on the edges and in the middle of the scanning region shows that the locking fluctuations are also smaller in the middle of the scanning range. The apparent spectral transition that appears in many of the spectra at 51 790 cm\(^{-1}\) is due to atmospheric absorption by \( \text{O}_2 \) in the laser’s oscillator which at this wavelength causes the laser to operate completely broad band. The signal rises approximately to the level present on the edges of the scanning range and then returns to the locked mode when the atmospheric \( \text{O}_2 \) absorption is passed by. Although there are four atmospheric absorptions in the laser scanning range, this effect is only observed at 51 790 \( \text{cm}^{-1} \) where the output of the oscillator is already fairly weak.

**FIG. 1.** Experimental apparatus with modified ArF laser: Grating (G), prisms (P\(_1\), P\(_2\)), windows (W\(_1\), W\(_2\), W\(_3\), W\(_4\)), slits (S\(_1\), S\(_2\)), mirrors (M\(_1\), M\(_2\)), output couplers (C\(_1\), C\(_2\)), cylindrical lenses (CL\(_1\), CL\(_2\)), back reflector (BR), Detection lenses (DL\(_1\), DL\(_2\)), image intensifier (II).
$J = N + 1$, and the blue shifted line contains both the $F_2$, $\bar{J} = \bar{N}$, and $F_3$, $\bar{J} = \bar{N}-1$, components. In Hund's case B, all three components have the same absorption probability, neglecting very small differences in the Hoenl–London factors and deviations for the lowest rotational states. Emission from the different components of the triplet occurs at almost the same wavelength and there is consequently no discrimination between the components due to the monochromator. In addition, the polarization of the emission coming from the three components is negligibly different even for the lowest $J$ states. It is therefore immediately clear that the three do not predissociate with the same probability. Empirically, one can conclude that the sum of the $F_2$ and $F_3$ lifetimes must be about equal the $F_1$ lifetime.

In Fig. 3, the triplet is fully resolved in several ($B \nu' = 15,16 - \nu'' = 3$ transitions due to the larger spin–spin and spin–rotation constants for $\nu' = 15$ and $\nu' = 16$. Note particularly the well isolated $\nu' = 16 - \nu'' = 3; R(19)$ triplet at 51 750 cm$^{-1}$. Although a quantitative simulation of this spectrum is elusive due to its increased complexity, the qualitative observation that $F_1$ states predissociate approximately half as quickly as $F_2$ and $F_3$ states is clear for $\nu' = 14,15$, and $16$.

The relative lifetimes of the triplet are directly related to the symmetry of the repulsive state involved, as will be shown in more detail below. From the relative coupling matrix elements calculated by Julienne$^6$ one can conclude that $\nu' = 14,15$, and $16$ are predissociated primarily by a state of $^3\Sigma_u^+$ or $^3\Pi_u$ symmetry, whose triplet lifetime patterns are identical. While a $^3\Pi_u$ state has been predicted to be important for predissociation of low vibrational levels, no potential calculations have suggested that a curve of this symmetry could predissociate $\nu' = 14,15$, or $16$. However, a $^3\Sigma_u^+$ curve has been calculated to lie in the correct energy range and to have a fairly large spin–orbit coupling magnitude. The assumption of $^3\Pi_u$ or $^3\Pi_u$, the latter thought to be dominant from absorption spectra, implies completely different triplet patterns. For example in the case of the well isolated $\nu' = 16 - \nu'' = 3; R(19)$ transition, the $F_1:F_2:F_3$ ratio's are 0.7:1.0:0.6 and 1.0:0.9:0.9, respectively, for $^3\Pi_u$ and $^3\Sigma_u^+$. This shows the sensitivity of measuring spin-state dependent lifetimes as opposed to rotation vibration dependent lifetimes.

The observed importance of the $^3\Sigma_u^+$ curve urged us to investigate the $\nu' = 11$ level. Based on the interpretation of absorption spectra, this vibrational state should be where the $^3\Sigma_u^+$ repulsive curve has the largest influence observable in our experiment (although $^3\Pi_u$ was still thought to be dominant for this level). Moreover, the $\nu' = 11$ state predissociates some 10 to 20 times faster than $\nu' = 14,15,16$ and rep-

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**FIG. 2.** Laser induced fluorescence spectrum: $\nu' = 14,15,16 - \nu'' = 3$ simulated (above), measured (below).

**FIG. 3.** Laser induced fluorescence spectrum: $\nu' = 15,16,17 - \nu'' = 3$ simulated (above), measured (below).
resents a much more important O-atom production channel in the atmosphere. In an effort to achieve a clear picture of the predissociation mechanism of \( v' = 11 \), the laser was operated with a resolution of 0.5 cm\(^{-1}\) by fully illuminating the oscillator's grating and excitation spectra of well isolated \( v' = 11 - v'' = 2 \) triplets were measured.

Figure 4 shows the excitation spectrum of the \( R(27) \) triplet. The Lorentzian influence on the line shape is clearly visible, note the long wings on the lines. Three simulated spectra are also shown based on the state symmetry dependence of the relative coupling matrix elements from Ref. 5. Only a \( ^3\Sigma_u^+ \) repulsive state reproduces the observed line shape. Therefore, it is clear that it is the dominant mechanism. This is in stark contrast to theoretical predictions which have considered \( ^3\Pi_u \) to be relatively unimportant. How LIF measurements of spin-state dependent lifetimes can lead to such different results than analysis of absorption spectra will be discussed below.

**IV. DISPERSION SPECTRA**

Figure 5 shows two dispersed fluorescence spectra. The lower spectrum results from broad band excitation where the oscillator of the laser is blocked. The broad emission lines come from excitation of all accessible molecular transitions within the gain profile of the laser and subsequent emission from a large number of rotational states in \( \text{O}_2 \): \( v' = 10, 11, 12, 14, 15, 16, \) and \( 17 \) as well as a small amount of \( \text{NO} \) \( D \)-state emission at 208.9 nm.\(^{11}\) Narrow band excitation of two coincident \( \text{O}_2 \) rovibronic transitions \( [v' = 10 - v'' = 2:R(29)] \) and \( v' = 16 - v'' = 3:R_1(19) \), Fig. 5 (above), produces single \( P \) and \( R \) emission lines, whose width is limited by the resolution of the monochromator and completely suppresses

![Diagram](image1.png)

**FIG. 4.** High resolution laser excitation spectrum of \( v' = 11 - v'' = 2:R(27) \) triplet. Simulated spectra assuming three different symmetries for the repulsive states.

![Diagram](image2.png)

**FIG. 5.** Dispersion spectrum resulting from broad band laser excitation (below) and from narrow band excitation (above). Vibrational states involved in narrow band excitation are noted at far right and final vibrational states after emission are listed below the emission lines. Note small amount of \( \text{NO} \) emission in broad band spectrum.
the NO emission. This shows clearly the absence of collisional effects even at atmospheric pressure in the flame since collisional relaxation of the B state would destroy the single P, R line emission observed. At longer wavelengths the O₂ emission lines split into two components, corresponding to the v" = 2 and 3 excitations. The emission resulting from v" = 2 excitation is always the red shifted component.

Because of the absence of quenching, very simple emission spectra arising from laser prepared single rovibronic levels are observed. Due to the large difference between the equilibrium bond lengths of the two electronic states, emission is observed to states as high as v" = 35. Data such as that in Fig. 5 (above) were analyzed to obtain relative emission probabilities to states within 0.3 eV of X-state dissociation. Great care was also taken to filter out UV emission that can appear in the monochromator's second order. The results are collected in Fig. 6.

These relative Einstein B coefficients are determined by factoring out the v' dependence for spontaneous emission as well as the overall experimental sensitivity as a function of the sum of the Franck–Condon factors for emission from one v' state to all v" states must be unity. The Franck–Condon factor sum for v' = 10 → v" = 0–23, for example, is only 0.28.12 This implies that the Franck–Condon factors to states between v" = 23 and dissociation sum to 0.62. The experimentally derived emission probabilities in this region (shown in Fig. 6) only account for about half of this value and are therefore clearly influenced by the internuclear separation dependent transition dipole moment and cannot be simply interpreted as Franck–Condon factors.

For higher v" states and consequently larger internuclear separations, transition dipole moments are not available but the variation with internuclear separation in this region is expected to be large for two reasons. First, in the separated atom limit the transition is forbidden so the transition dipole moment is going asymptotically to zero. Second, the sum of the Franck–Condon factors for emission from one v' state to all v" states must be unity. The Franck–Condon factor sum for v' = 10 → v" = 0–23, for example, is only 0.28.12 This implies that the Franck–Condon factors to states between v" = 23 and dissociation sum to 0.62. The experimentally derived emission probabilities in this region (shown in Fig. 6) only account for about half of this value and are therefore clearly influenced by the internuclear separation dependent transition dipole moment and cannot be simply interpreted as Franck–Condon factors.

In general, the emission probabilities to these high vibrational states are strikingly large, especially for emission from v' = 10, 11. Emission from v' = 14, 15, and 16, to high vibrational states is smaller as the transition dipole moment is approaching the forbidden free-atom limit (O' D + O' D → O' P + O' P).

The resolution of P and R emission allows the rough determination of the rotational B constants for these high vibrational states. Figure 7 shows the results of this analysis with a comparison to already known rotational constants.14 While the scatter in the data is relatively large, the tendency toward ever smaller B values is consistent with the lengthening of the chemical bond at higher vibrational quantum numbers and further confirmation of the correct assignment of the observed emission.

V. DISCUSSION

A. Spin-state selective predissociation in O₂

The interpretation of spin-state selective predissociation in the O₂ B state depends on the spin–orbit coupling of the B'2Σ⁺ state to curves of differing symmetry which depends only on known spin–spin and spin–rotation splitting constants. In Hund's case A notation, such transitions must conserve the Ω quantum number (the projection of total electronic angular momentum) and parity. The critical matrix elements take the following form.

$$|\langle F_i | L · S | \text{REPELATIVE}, \Omega, \pm \rangle|^2,$$

where: F_i are the B-state triplet components, ± is the parity of the repulsive curve. The F_i wave functions of the B state can be expressed in Hund's case A wave functions as follows.
B. The extension of the \( \text{O}_2 \) \( X \)-state PES

When possible diatomic PES's should be determined from spectroscopically derived molecular structure constants. Such constants for the \( X \) state of \( \text{O}_2 \) have been determined with reasonable accuracy up to \( v'' = 28 \).\(^1\) From such information Franck–Condon factors up to \( v'' = 23 \) have been derived\(^12\) and the agreement with the results of this work is an indirect confirmation of the accuracy of the \( X \) state PES up to \( v'' = 23 \). However, molecular constants of higher vibrational states are not yet available and the region of the PES between 3.7 (\( v'' = 23 \)) and 5.1 eV (dissociation) remains undetermined. It is precisely this portion of the PES that plays a crucial role in the chemistry of \( \text{O}_2 \), unimolecular dissociation and three-body recombination being the most pertinent examples. We propose that the relative emission information derived from this experiment can be used to extract the PES in this energy region.

To achieve this, one must vary an empirical \( X \)-state PES, calculate the Franck–Condon integrals, the dipole-moment integrals, and check against the observed emission probabilities. One must further check that the PES delivers \( B \) constants that agree with those derived from the \( P,R \) emission lines of this experiment, Fig. 7. Normally, this would be a very unsatisfactory method for PES determinations, because the integrals mentioned are quantities averaged over all parts of the PES. However, the Franck–Condon factors between high vibrational levels of the two electronic states result mainly from overlap of tightly localized wave functions at the outer turning point. This suggests that such data could be quite sensitive to the PES in the region of dissociation. The attractive part of the \( B \)-state PES is already relatively well determined up to \( v' = 15 \). We hope that the results of this experiment will stimulate further work in this direction.

VI. CONCLUSIONS

Specific new results on the spectroscopy and photodissociation dynamics of \( \text{O}_2 \) have been obtained by measuring LIF spectra with a tunable ArF laser. The spin-state selective predissociation has been observed in \( v' = 11,14,15, \) and 16 of the \( B \)-state providing a clear picture of the mechanism of \( \text{O}_2 \) predissociation for these states. The results show that the spin-state dependence of the predissociation lifetimes is a much more sensitive probe of the identities of the repulsive curves than is the vibration–rotation state dependence derived from absorption spectra. It is found that a repulsive \( ^3 \Sigma_u^+ \) state is important for high vibrational levels \( (v' = 14,15,16) \). Predissociation of \( v' = 11 \) by a \( ^3 \Pi \) repulsive state was also clearly observed. These results are in direct contrast to previous studies.

By preparing single rovibronic states in high vibrational levels, simple dispersion spectra have been obtained providing relative emission probabilities to \( X \)-state vibrational levels within 0.3 eV of dissociation. This data may provide the needed information to determine the \( \text{O}_2 \) PES out to the dissociation limit.


5(a) P. S. Julienne, J. Mol. Spectrosc. 63, 60 (1976); (b) P. S. Julienne and M. Krauss, ibid. 56, 270 (1975).


