

Spin torque on molecular rotation induced by polarized electrons

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We have measured the linear and circular polarization of 391.4 nm $N_2^+ B^2\Sigma_u^+(v'=0) \rightarrow X^2\Sigma_g^+(v''=0)$ P -branch fluorescence produced by spin-polarized electron impact excitation of $N_2 X^1\Sigma_g^+(v=0)$ ground states. The resulting nonzero values of the circular polarization indicate that spin-rotation coupling, in conjunction with electron exchange excitation, acts to produce oriented rotational angular momenta in the excited molecular state prior to decay.

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I. INTRODUCTION

Use of spin-polarized electrons in collisions with atoms and molecules provides a direct probe of spin-dependent collision dynamics [1,2]. A prototypical example of this is the production of circularly polarized fluorescence resulting from the transfer of spin angular momentum to orbital orientation in a collisions-excited atomic target [3]. In such reactions, the target becomes spin polarized by an exchange collision with a polarized electron beam. For the fluorescence to be circularly polarized, there must exist a mechanism to couple the newly oriented spin angular momentum, S , with the initially unoriented electron orbital angular momentum L . This is accomplished by the spin-orbit interaction, in which L and S couple to form J , which remains fixed in space (ignoring nuclear spin). Due to the precession of L and S about J , L becomes spatially oriented and circular polarization (relative Stokes parameter $P_3 \neq 0$) can be observed, provided that the time scale of the $L \cdot S$ coupling is less than or comparable to the excited state lifetime.

Molecular orientation can be considered in a similar manner. Recently, circular polarization of the fluorescence from excited H_2 triplet molecular states produced by spin-polarized electron impact has been observed [4]. (The identification of triplet states ensures that spin exchange has occurred.) Values of circular polarization normalized to the spin polarization of the electron beam, P_3/P_e , of ~ 0.1 were observed for a mixture of Fulcher band transitions in the 595–605 nm wavelength region. The mechanism for this polarization production is similar to that in atoms, but with an added layer of complication due to the molecular angular momentum coupling scheme (see Fig. 1). In Hund's case (b) molecules such as H_2 , the projection of the electronic orbital angular momentum onto the internuclear axis (Λ) couples with the nuclear rotational angular momentum (N) to form the total orbital angular momentum (K), which in turn couples with the electron spin (S) to form the total angular momentum (J). (Note that N is not a good quantum number unless $\Lambda=0$ so that $K=N$.) Circular polarization is produced when K is oriented by virtue of its coupling with S through a spin-orbit term in the molecular Hamiltonian $A(S \cdot K) = AS \cdot (\Lambda + N)$. As in atoms, this occurs over the spin-coupling time, i.e., the inverse precession frequency of S

about J . Such times are comparable to those in the constituent atomic cases. In the experiments of Green *et al.* [4], the fluorescence was primarily from Π states ($\Lambda=1$), and the ultimate molecular orientation was mainly due to the electron spin-orbit interaction with Λ (as opposed to N). This is because the magnetic field in the rest frame of the electrons due to their orbital motion is much larger than that due to the (relatively slow) rotation of the nuclei themselves.

In the experiment reported here, we demonstrate that the collision-induced orientation of K , and hence P_3 production, is possible even in the absence of electronic orbital angular momentum along the internuclear axis ($\Lambda=0$ and $K=N$). The mechanisms responsible for this torque on the molecular nuclei are the direct coupling of S to the magnetic field produced by the "current loop" of nuclear rotation, as well as the coupling of S to the residual electronic orbital angular momentum perpendicular to the internuclear axis, which also produces a magnetic field along K [5–7]. [The latter mechanism can also be attributed to nuclear rotation, however, in that it requires uncoupling of the total electron orbital angular momentum (L) from the internuclear axis.] To date, the

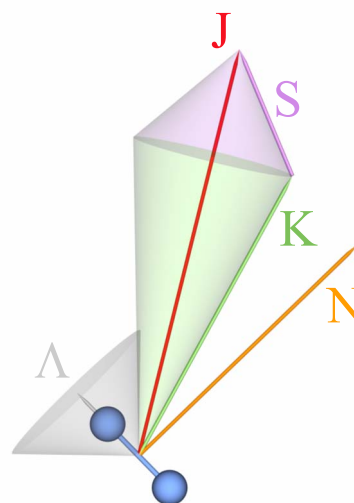


FIG. 1. (Color online) Schematic of Hund's case (b) coupling for a $\Lambda=1$ (Π) state. The S and K vectors precess around the total angular momentum J in the fine-structure coupling time. Initial orientation of S leads to orientation of J and K . The vector N is always perpendicular to the internuclear axis, and the precession loci of L , K , and S are indicated by transparent cones.

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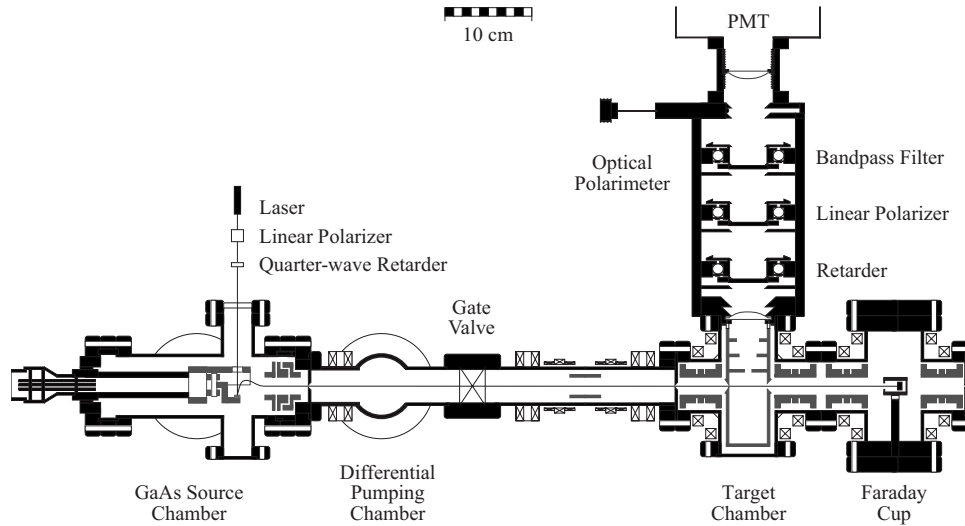


FIG. 2. Experimental apparatus.

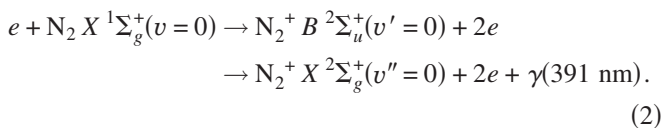
effects of these rotational couplings have only been observed as “spin doubling” in molecular spectra [8], although it has been discussed as a possible mechanism for spin relaxation in triplet alkali dimers [9,10]. For ${}^2\Sigma$ states, this so-called “spin-rotation” term in the Hamiltonian, $\gamma(\mathbf{K}\cdot\mathbf{S})$, gives rise to energy splittings

$$\Delta E_{\text{SR}} \approx \gamma\left(K + \frac{1}{2}\right). \quad (1)$$

Our experiment provides direct evidence for molecular orientation due to spin-rotation coupling in electron-molecule collisions.

II. EXPERIMENT

We have studied the excitation-ionization reaction with N_2 excited by spin-polarized electrons leading to fluorescence in the first negative system of N_2^+ ,



Our experimental apparatus is shown in Fig. 2. To obtain a beam of spin-polarized electrons, circularly polarized 785 nm light from a laser diode irradiates an unstrained GaAs crystal. Layers of cesium and oxygen are applied to the crystal in order to produce a negative electron affinity surface, allowing electrons to escape [11] with a typical yield of 1–10 $\mu\text{A}/\text{mW}$ and an energy width <0.4 eV. The crystal is held at a potential of -250 V to define the beam transport energy. Photoelectrons are electrostatically deflected by 90° so that the spin polarization becomes transverse with respect to the beam direction. The spin direction is determined by rotating the quarter-wave retarder controlling the laser polarization. The amount of spin-polarization, P_e , is 27%, as measured by optical electron polarimetry [12]. A differential pumping region separates the field-free target gas cell from the source. The target potential is varied to change the interaction energy. Upon exiting the target, the electron beam is

collected in a Faraday cup for current normalization. A lens above the target cell images light from excited molecules into an optical polarimeter [13], comprising a rotatable quarter-wave retarder, linear polarizer, bandpass filter and Hamamatsu R946 photomultiplier tube.

Data acquisition consisted of incrementing the beam energy every 5 s; for each energy interval, the integrated counts, average current, and average pressure were measured. After a single energy scan was completed, the beam energy was reset to the starting value and the angle of the retarder in the optical polarimeter incremented by 22.5° . This process continued until a full rotation of the polarimeter retarder was achieved. The angle of the laser quarter-wave retarder was then incremented by 90° , changing the spin direction of the electrons to help minimize instrumental asymmetries. A single “run” comprised several full rotations of the laser retarder.

Below-threshold unnormalized count rates were averaged to determine the background, which was then subtracted from all count rates. (Chauvenet’s criterion was applied to remove anomalous count rates [14].) Above-threshold count rates were then normalized to pressure and current, and count rates corresponding to unique energies and orientations of the optical retarders were combined to give final intensities and their uncertainties. For a given energy, the intensities for different optical orientations were subjected to an inverse Fourier transform to obtain Fourier coefficients related to the desired Stokes parameters [15].

The polarization of molecular fluorescence depends on the rotational transitions observed. To illustrate our optical sensitivity, Fig. 3 shows a spectrum for the $\text{N}_2^+ B^2\Sigma_u^+(v'=0) \rightarrow X^2\Sigma_g^+(v''=0)$ band excited by electron bombardment [16]. As expected, there is a 2:1 intensity alternation of even K'' vs odd K'' rotational transitions due to the nuclear spin statistics of ${}^{14}\text{N}_2$ [8]. The effect of the narrow bandpass filter [center wavelength 391.6 nm; full width at half maximum (FWHM) 0.92 nm] used for our experiment is shown in the lower plot, indicating that our data is due only to P -branch ($K' - K'' = -1$) fluorescence, the R branch ($K' - K'' = +1$) being almost completely suppressed. Also, we preferentially

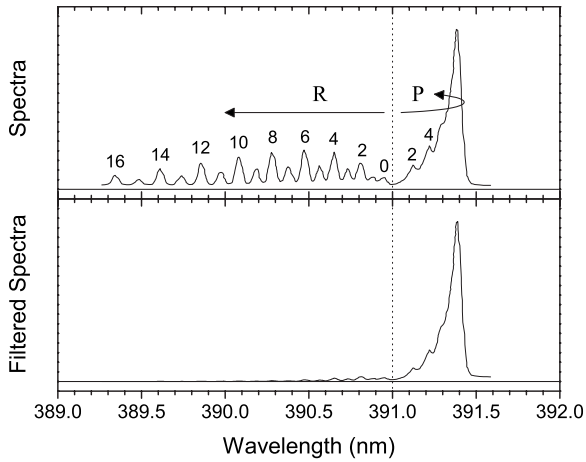


FIG. 3. Upper figure, rotational spectrum for the 391.4 nm band taken from Ref. [16]; some final rotational states for the $P(K'')$ and $R(K'')$ branches are labeled. Lower figure, spectrum modified by our bandpass filter transmission.

detect transitions near the band head $P(K'' \sim 14)$. No significant contaminant molecular or atomic lines are expected to coexist in the profile of the bandpass filter. A transition in the N_2 Goldstein-Kaplan $C' \ ^3\Pi_u(v'=0) \rightarrow B \ ^3\Pi_g(v''=8)$ system at 391.5 nm exists, but it is apparently much weaker than the transition under study, as we observe negligible intensity from this band below the threshold for $B \ ^2\Sigma_u^+$ state production.

III. RESULTS

Our linear and circular polarization data presented in Fig. 4 were acquired at a pressure of 4×10^{-2} Pa to avoid potential effects of radiation trapping. Cascade contributions from higher-lying states are not expected [17]. Our P_1 values are consistent with those of Ref. [18], but we note that the interference filter used in that work transmitted both the P - and most of the R -branch fluorescence. It is difficult to assess what the kinematically required threshold value of P_1 should be. For production of the $B \ ^2\Sigma$ state, preferential excitation of $M_J = \pm \frac{1}{2}$ levels is expected [19]. This would yield positive P_1 values for the thermal ensemble of rotational states present in a room-temperature target, which we do observe. In addition, the effects of hyperfine and fine-structure depolarization are significant [20], leading to an overall reduction in the degree of linear polarization at all energies. The measured values of P_2 (the $45^\circ/135^\circ$ linear polarization fraction), are consistent with zero. This is expected, as the total spin of the collision system should be a good quantum number due to the low Z of the target nuclei [21].

Since the molecular states of interest are all of Σ character, the angular momentum coupling is described by Hund's case (b). Figure 5 depicts this coupling scheme, wherein the vectors \mathbf{K} and \mathbf{S} couple to form the spatially fixed \mathbf{J} (total angular momentum excluding nuclear spin). Vectors \mathbf{K} and \mathbf{S} precess about the spatially fixed \mathbf{J} with a period of < 0.1 ns ($K=1$) corresponding to the $\gamma(\mathbf{K} \cdot \mathbf{S})$ "spin-doubling" term in the molecular Hamiltonian [22]. This spin-rotation coupling

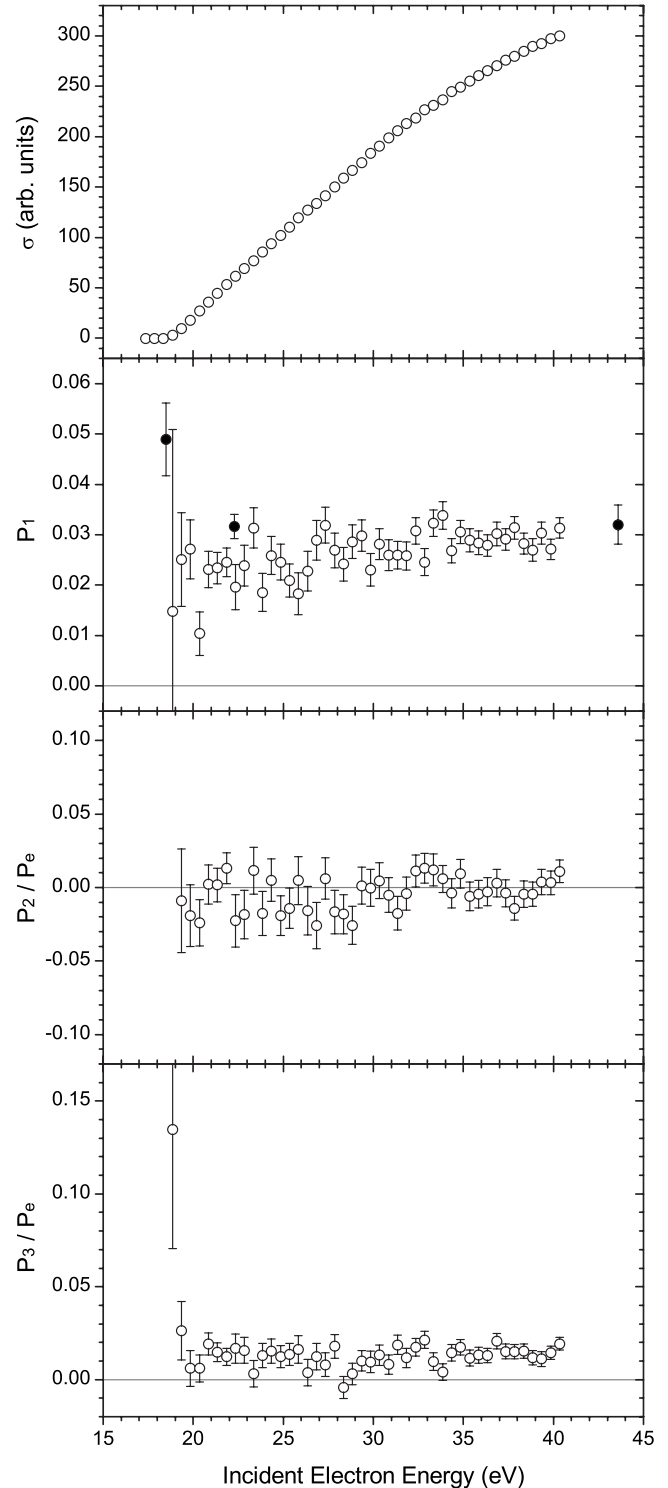


FIG. 4. Relative optical emission cross section and fluorescence polarizations vs electron energy for the 391.4 nm $N_2^+ B \ ^2\Sigma_u^+(v'=0) \rightarrow X \ ^2\Sigma_g^+(v''=0)$ band. Data of this work is represented by open circles (1σ errors); data of Ref. [18] by filled circles (errors represent 98% confidence level). Threshold is at 18.75 eV [23].

time scale is much less than the lifetime of the $B \ ^2\Sigma_u^+$ system (61 ns [24]), so time-averaged orientation of \mathbf{K} can develop.

The nonzero value of P_3/P_e for the 391.4 nm fluorescence provides direct evidence of molecular orientation due

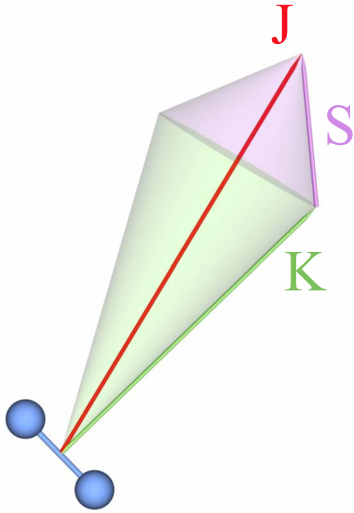


FIG. 5. (Color online) Schematic of Hund's case (b) coupling for a $\Lambda=0$ (Σ) state. The S and K vectors precess around the total angular momentum J . Initial orientation of S leads to orientation of J and K . The vector K is always perpendicular to the internuclear axis.

to this coupling. Fitting a constant to the data gives an energy-averaged value of 0.0133(8), with a reduced $\chi^2 = 1.04$. However, the observed values are relatively small when compared to the circular polarizations detected for atomic and even molecular H_2 triplet-state fluorescence. There are several reasons that may explain the small values of circular polarization we observe. The first is related to the fact that the B state is a doublet, so electron exchange is not guaranteed, and the initial spin orientation may be diluted relative to a triplet state by an exchange-to-direct excitation cross-section ratio. Another factor to consider is the relative magnitude of rotational state angular momenta in N_2 vs H_2 . For larger values of K , spin orientation is less effective at producing appreciable nuclear orbital orientation, resulting in smaller P_3 values. Room-temperature rotational states in nitrogen are populated up to $K \sim 20$, while for hydrogen the maximum K is ~ 3 . Thus the doublet state spin momentum of $\frac{1}{2}\hbar$ is small compared with the rotational angular momen-

tum of $\sqrt{K(K+1)}\hbar$. In fact, this ratio ($\sim \frac{1}{40}$ for maximum K) is roughly an order of magnitude less than that for triplet state H_2 ($\sim \frac{1}{3}$ for maximum K). This effect alone could account for the differences in the observed P_3/P_e from these two molecular species. Finally, the results are susceptible to both fine and hyperfine depolarization [20,25]. Typical hyperfine splittings in the excited states of atomic nitrogen are of the order of 10^8 Hz [26], so molecular depolarization should occur in ~ 10 ns.

In this regard, we mention an earlier experiment performed by the Münster group in which the circular polarization for the $N_2 C^3\Pi_u(v'=0) \rightarrow B^3\Pi_g(v''=0)$ 337.1 nm band (second positive system) excited by spin-polarized electrons was measured. Values of P_3/P_e were found to be zero within an uncertainty of 2×10^{-3} [27]. This result is surprising, given that the fully oriented (triplet state) electrons couple strongly to the internuclear axis [Hunds case (a)] and should provide orientation of K . We have since confirmed this null result for the $N_2 C^3\Pi_u(v'=0) \rightarrow B^3\Pi_g(v''=2)$ 380.5 nm band. We speculate that it is due to the fact that in both experiments there was no resolution of the rotational band structure of the upper state, giving both P - and R -band fluorescence equal weight. This would tend to cause the integrated value of P_3 to average to zero [28]. Further rotationally isolated measurements are desirable in order to confirm this conjecture.

IV. CONCLUSION

In conclusion, we have shown that orientation of molecular states can occur via spin exchange even in the absence of direct spin-electron orbit coupling. This results from the magnetic coupling of the electron spin with both nuclear rotation directly, and with its own residual orbital angular momentum perpendicular to the internuclear axis. This experiment illustrates the usefulness of spin-polarized electrons as a probe of molecular collision dynamics.

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