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UV Photodissociation Dynamics of HN₃ in 190-248 nm

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The H+N₃ channel in the ultraviolet photodissociation of HN₃ has been investigated from 190 nm to 248 nm using the high-n Rydberg H-atom time-of-flight technique. Product translational energy distributions as well as product angular anisotropy parameters were determined for the H+N₃ channel at different photolysis wavelengths. N₃ vibrational state distribution has also been derived from the product translational energy distribution at these wavelengths. Above photolysis wavelength 225 nm, HN₃ predominantly dissociate through the repulsive state. Below 225 nm, a new slow channel starts to appear at 220 nm in addition to the existing channel. This channel is attributed to a ring closure dissociation channel to produce the cyclic N₃ product. As photolysis energy increases, this new channel becomes more important.

Key words: Photodissociation dynamics, H atom channel, UV photolysis, Product vibrational state distribution

I. INTRODUCTION

The absorption spectrum of HN₃ is composed of at least three absorption systems between 180-300 nm: a weak one peaking at 264.3 nm and two strong ones peaking at 204 and 190 nm, respectively [1-4]. These absorption features are broad and nearly structureless. The weak feature at λ>220 nm has been assigned to AA′′−X'A′ [2,3,5]. At shorter wavelengths, the BA' and CA'′ electronic states are optically accessible, and the resulting UV absorption spectrum is strong, but no discrete structure is present.

It is known that the UV photodissociation of HN₃ occurs primarily via two spin-allowed channels [1-16]:

\[
\text{HN}_3(XA') \rightarrow \text{H}(\tilde{2}S) + \text{N}_3(X^2\Pi_g) \rightarrow \text{NH}(a^1\Delta) + \text{N}_2(X^1\Sigma_g^+) \tag{1}
\]

The NH+N₂ channel has been extensively studied at several different wavelengths [4,6-11]. NH is produced in the lowest singlet \((a^1\Delta)\) instead of the \(X^3\Sigma^-\) ground state because of the spin symmetry rule. The H-atom has been investigated by Comes and coworkers by laser induced fluorescence method [12,13], and H-ion TOF spectra [14]. The average center-of-mass translation energy \((E_{\text{trans}})\) release has been found to be 68% of the total available energy at 248 nm. The H atom quantum yield has been measured at several wavelengths: \(\phi_{266}=0.04, \phi_{248}=0.20\). At 266 and 248 nm, the product anisotropy parameter is close to -1, indicating a perpendicular electronic transition \((AA''\rightarrow XA')\). Using the H-ion TOF spectra method [14], H−N bond energy is determined to be: \(D_0(H-N_3)=30850\pm400\) cm⁻¹. The main structures observed in the distribution were assigned to a progression in the \(v_1\) symmetric stretching mode of N₃ and the relative population of the \((0,0,0):(1,0,0):(2,0,0):(3,0,0):(4,0,0)\) modes was found to be 0.09:0.19:0.28:0.29:0.15. Contributions to the TOF profile from other active vibrations could not be ruled out, because the resulting H+ TOF profile was poorly resolved.

Recently, the high-n Rydberg H-atom time-of-flight (TOF) technique has also been employed to investigate the H-atom channel. Cook et al. reported the translational energy spectra of the H+ N₃(X) products following excitation at eight different excitation wavelengths (λ≥240 nm) [15]. Based on their experimental results and analysis, they concluded that the N₃ fragments are formed in a progression of levels involving both the symmetric stretch mode \((v_1,0,0)\) and the symmetric stretch mode combined with one and two quanta of bending motion, \(v_2\). The rotational energy disposal in the N₃(X) products was also estimated using an impulsive dissociation model, suggesting that the most probable impact parameter \(b\) is 1.26±0.05 Å. The dissociation energy of the H−N₃ bond is determined to be: \(D_0(H-N_3)=30970\pm50\) cm⁻¹. The \(N_3(X)\) vibrational state population distributions are determined at eight photolysis wavelengths. Zhang et al. have also used Rydberg H atom TOF method to investigate the HN₃ photodissociation at 248.3 and 193.3 nm [16]. At 248.3 nm, Zhang et al. assigned the vibrational structures in the product translational energy spectra, similar to Ref. [15]. The H−N₃ bond energy was determined to be \(D_0(H-N_3)=31020\pm180\) cm⁻¹. At 193.3 nm, they...
found that the product anisotropy parameter is clearly energy-dependent, implying multiple dissociation pathways are responsible.

The spectroscopy of N\textsubscript{3} is rather complicated due to several interactions (Renner-Teller and spin-orbit splitting, etc.) [17-20]. Early studies demonstrated an absorption band which was attributed to an electronic transition at \( \sim \)270 nm [17]. The ground state exhibits a spin-orbit splitting of \( A_0,\text{eff} = -71.26 \text{ cm}^{-1} \), and a Renner-Teller splitting of \( \varepsilon \omega_2 = -94.38 \text{ cm}^{-1} \) for the bending vibration. Subsequent wavelength dispersed LIF studies [18] yielded approximate values for the ground state symmetric stretch and bending vibrational frequencies of \( \omega_1 = 1320 \text{ cm}^{-1} \) and \( \omega_2 = 457 \text{ cm}^{-1} \) respectively. The asymmetric stretch frequency \( (\omega_3 = 1644.678 \text{ cm}^{-1} [19]) \) has been studied by Fourier transform spectroscopy, and laser magnetic resonance [20]. Recent calculation [21] reported a linear symmetric ground state with two equal N-N bond lengths of 1.1818 Å in good agreement with the experimental value [19,20]. Meier et al.'s calculation showed that the HN\textsubscript{3}\( (A) \) state PES indicates a barrier (with an estimated barrier height \( \sim 2300 \text{ cm}^{-1} \)) along the H--N\textsubscript{3} dissociation coordinate.

In this work, we re-investigated the photodissociation dynamics of the HN\textsubscript{3} molecule at different wavelengths lower than 248 nm in an effort to study how the dynamics changes when the photolysis energy increases.

### II. EXPERIMENTS

The technique of H/D Rydberg atom TOF spectra technique has been described in detail elsewhere [22-24]. In this experiment, a skimmed, pulsed molecular beam of HN\textsubscript{3}, seeded in helium (mixing ratio \( \sim 2\%\), total pressure \( \sim 101 \text{ kPa} \)), is crossed perpendicularly with the photolysis laser beam. In the wavelength range of 225-280 nm, a frequency doubled dye laser (Sirah, PESC-G-24) pumped by a Nd:YAG (Spectra Physics Pro-290) is used. Laser wavelengths in the range of 188 nm to 225 nm are generated by frequency mixing of the second harmonic of the dye laser output with the Nd:YAG laser. Laser beams were then focused into a cell with Kr/Ar mixing gas where four wave mixing at 121.6 nm was generated. The 366 nm light was generated by frequency doubling a Radiant dye laser output (Jaguar, D900MA), operating at \( \sim 732 \text{ nm} \), which was pumped by a port of the 532 nm light of the same Nd:YAG laser to excite the H atoms from the n=2 level to a high-n Rydberg state below the ionization threshold. The two 121.6 and 366 nm laser beams were overlapped exactly in the interaction region both in space and in time. Ions and electrons formed in this region were extracted by a small electric field placed across the interaction region.

The tagged H atoms then fly away from the interaction region to reach the MCP detector. The Rydberg H atoms are efficiently ionized as soon as they pass a fine grounded metal grid in front of the MCP detector by the strong electric field applied to the front MCP plate. The total distance from the interaction region to the front face of the detector is \( \sim 333 \text{ mm} \). The signal obtained is amplified by a preamplifier, discriminated by a discriminator, sent to both a digital oscilloscope for visual display and a multi-channel scaler (P7888-2(E) FASTCOMTEC) for accumulation (typically more than 10\textsuperscript{7} laser shots per TOF profile) and subsequent data analysis.

HN\textsubscript{3} was prepared by heating sodium azide (NaN\textsubscript{3}) in excess stearic acid under vacuum for 3-4 h at 80-100 °C [24,25]. The HN\textsubscript{3} sample was stored in a stainless steel container and He gas was filled to produce 2\% HN\textsubscript{3}/He ratio (the total pressure made up to 101-404 kPa with pure helium). Purity was checked by mass spectrometry (SRS, RGA200).

### III. RESULTS AND DISCUSSION

#### A. Dissociation dynamics from 230 nm to 264 nm

The time-of-flight spectra (TOF) of the H-atom product from the photodissociation of HN\textsubscript{3} at many different wavelengths from 230 nm to 264.3 nm were measured using the method described above. These measurements were carried out with the rotating detector direction perpendicular and parallel to the photolysis laser polarization. TOF spectra at four specific wavelengths are shown in Fig.1. Since the photolysis laser was generated using a frequency doubling scheme, the photolysis laser polarization was fixed to parallel to the molecular beam direction for convenience in this wavelength region. The TOF spectra shown in Fig.1 were then converted to the total product translational energy \( (E_{\text{trans}}) \) distributions.

Since the total product translational energy \( (E_{\text{trans}}) \) deposited into the two products (H and N\textsubscript{3}) can be written as [16,26]

\[
E_{\text{trans}} = h\nu + E_{\text{int}}(\text{HN}_3) - D_0(\text{H} - \text{N}_3) - E_{\text{int}}(\text{N}_3)
\]

where \( h\nu \) is the photon energy of the photolysis laser,
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FIG. 1 TOF spectrum recorded at the photolysis wavelengths of (a) $\lambda=248$ nm, (b) $\lambda=240$ nm, (c) $\lambda=235$ nm, (d) $\lambda=230$ nm, with the MCP detector perpendicular to the laser polarization (solid lines) and parallel to the laser polarization (dotted lines).

$E_{\text{int}}$(HN$_3$) is the internal energy of the HN$_3$ molecule, $D_0$(H$-$N$_3$) is the dissociation energy of the H$-$N$_3$ bond and $E_{\text{int}}$(N$_3$) is the internal energy of the N$_3$ product. HN$_3$ is cooled down to very low temperature in the supersonic expansion beam, therefore $E_{\text{int}}$(HN$_3$)$\approx$0. $E_{\text{trans}}$ can then be simplified to:

$$E_{\text{trans}} = h\nu - D_0(\text{H} - \text{N}_3) - E_{\text{int}}(\text{N}_3) \quad (4)$$

The total product translational energy distribution is also the internal energy distribution of the N$_3$ product. Thus, the structures observed in the product translational energy distribution are all due to the ro-vibrationally excited N$_3$ products. Here the photolysis energy, $h\nu$, is known exactly. From Eq.(4), if the maximum translational energy limit ($E_{\text{MAX}}$) in the product translational energy distribution corresponds to the N$_3$ ground ro-vibrational state, $D_0$(H$-$N$_3$) should be equal to $h\nu - E_{\text{MAX}}$. Using the extrapolated onset of $E_{\text{MAX}}$ in Fig.2, the H$-$N$_3$ bond energy, $D_0$(H$-$N$_3$) is determined to be 30910±100 cm$^{-1}$. This result is consistent with the value obtained by Cook et al., $D_0$(H$-$N$_3$)$=30970\pm50$ cm$^{-1}$ [15] and with the value by Zhang et al., $D_0$(H$-$N$_3$)$=31020\pm180$ cm$^{-1}$ [16].

The vibrational structures observed in all the translational energy distribution at all photolysis wavelengths studied here show a clear progression of doublets. These structures, given the availability of accurate theoretical vibrational term values [21], could be assigned to a progression in the N$_3$ symmetric stretching mode (characteristic spacing $\approx$1320 cm$^{-1}$), $(v_1,0,0)$, together with a progression of the symmetric stretching mode with one quantum of bending motion, $(v_1,1,0)$, as did in previous similar works by Cook et al. [15] and Zhang et al. [16].

In order to understand how the nascent N$_3$ product vibrational state distribution changes with the photolysis energy, we have also carried out a wavelength dependence study of the HN$_3$ photodissociation. Figure 3 shows an experimental translational energy spectrum obtained at 235 nm. In the simulation, a full width half maximum (FWHM) of 400 cm$^{-1}$ is used. In one of the simulation, we have included (lower panel in Fig.3) only progressions of the $(v_1,0,0)$ and $(v_1,1,0)$ vibrational series assuming only $v_2=0,1$ vibrational states are excited. Clearly, it is hard to simulate satisfactorily the translational energy spectrum. However, by including the series of $(v_1,2,0)$, the fitting to the translational energy spectrum is much improved (middle panel in Fig.3). This suggests that the N$_3$ products with $v_2=2$ $(v_1,2,0)$ vibrationally excitation are also likely present. By including the three series of the vibrational states, we have successfully simulated all the translational en-

FIG. 2 The product translational energy ($E_{\text{trans}}$) distribution derived from the experimentally measured H atom TOF spectra recorded at (a) $\lambda=248$ nm, (b) $\lambda=240$ nm, (c) $\lambda=235$ nm, (d) $\lambda=230$ nm, with $\theta=90^\circ$ (dotted lines), 0$^\circ$ (solid lines).

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FIG. 3 Experimentally derived internal energy profiles together with simulations for photolysis of HN$_3$ at $\lambda$=235 nm. (a) Individual contributions of N$_3$ vibrational state products; (b) includes progressions in the ($v_1$,0,0), ($v_1$,1,0) and ($v_1$,2,0) modes; (c) only progressions in the ($v_1$,0,0) and ($v_1$,1,0) modes are included.

FIG. 4 Relative populations of all clearly resolved N$_3$(X) vibrational states at four different photolysis wavelengths were obtained for both ($v_1$,0,0) and ($v_1$,1,0) series. Here, we normalized the population of the first vibrational state to 1.

energy spectra at the photolysis wavelengths from 225 nm to 254 nm. In the translational energy spectrum shown in Fig.3, most of the N$_3$ vibrational structures are not well resolved. However, the peak at $\sim$7900 cm$^{-1}$ in the translational energy distribution at the photolysis wavelength of 235 nm shown in Fig.3, which can be clearly assigned to the (5,2,0) vibrational state, strongly supports the argument that the ($v_1$,2,0) vibrationally excited N$_3$ products are present.

From the translational energy distributions shown in Fig.2, the average transition energy ($\langle E_{\text{trans}} \rangle$) at 230, 235, 240, and 248 nm are determined to be 64.6%, 66.5%, 68%, and 73%, respectively. The translational energy percentage decreases as photolysis energy decreases. The relative populations of ($v_1$,0,0) and ($v_1$,1,0) ($v_1 \leq 6$) different vibration states were also determined and shown in Fig.4. In this figure, the relative population of the lowest vibrational state at each photolysis wavelength is normalized to one. Clearly, the vibrational excitation of the $v_1$ mode increases as the photolysis energy increases. As photolysis energy increases, the $v_2$=1 state population also increases relative the $v_2$=1 state. This indicates the vibrational excitation of the N$_3$ radical increases in both vibrational modes that are excited. It is interesting to point out, however, that the rotational excitation of the N$_3$ product remains almost the same since the peak width in the translational energy distributions are similar in this photolysis energy regime.

Since product translational energy distributions, or N$_3$ product internal energy distribution can be obtained, were determined at two different detection directions that are perpendicular or parallel to the photolysis laser polarizations in this experiment, product angular anisotropy parameters can also be derived.

Figure 5(a) shows the translational energy dependent product anisotropy parameter obtained for the photolysis energy of 240 nm. Between the translational energy range of 2000-11000 cm$^{-1}$, the anisotropy parameter is between $-0.7$ to $-0.8$, and is only weakly dependent on the translational energy. It is noted that anisotropy parameters below the translational energy of 2000 cm$^{-1}$ is not provided because no signal is observed in this range at the photolysis wavelengths described here. Anisotropy parameter at other photolysis wavelengths in this photolysis energy region shows similar behaviors. The overall anisotropy parameter at different photolysis wavelengths in this regime only varies slightly.

Figure 6 shows the overall angular anisotropy parameters at the photolysis wavelengths of 264.3 nm to 188 nm, in which the overall anisotropy parameters at the photolysis wavelengths from 264.3 nm to 230 nm are all around $-0.7$. These results suggest that the photodissociation dynamics of HN$_3$ in this photolysis energy regime is similar, and the dissociation occurs directly via the excited potential energy surface to produce a linear N$_3$ radical product. This conclusion is quite consistent with previous experimental results by Cook et al. [15] and Zhang et al. [16].

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FIG. 5 Anisotropy parameter $\beta$ as a function of the total transitional energy ($E_{\text{trans}}$) at three typical photolysis wavelengths: (a) $\lambda=240$ nm, (b) $\lambda=217$ nm, and (c) $\lambda=193.3$ nm.

FIG. 6 The photolysis wavelength dependence of the overall angular anisotropy parameter $\beta$.

B. Dynamics from 225 nm to 213 nm

Figure 7 shows that the TOF spectra of the H-atom product obtained at the photolysis wavelengths of 225, 220, 217, and 213 nm with the detector at both the perpendicular and parallel directions. Since photolysis laser wavelengths below 225 nm was generated using a frequency mixing scheme as described above, the photolysis laser polarization was fixed to be perpendicular to the molecular beam direction for convenience in this wavelength region. The TOF spectra were converted to the product translational energy distributions, and are shown in Fig.8. From this figure, the dynamical feature in the translational energy distribution starts to become a little congested and the vibrational structures eventually disappear all together at the 213 nm photodissociation. This is likely due to the larger rotational excitation of the N$_3$ product because more available energy is deposited into the internal degrees of freedom of the N$_3$ radical product. Furthermore, as the photolysis energy increases, the overall product angular anisotropy parameter decreases noticeably. From Fig.6, the overall anisotropy parameter decreases from $-0.6$ to $-0.1$ in this wavelength regime, suggesting the dynamics is significantly altered in this range. From the translational energy distributions shown in Fig.7, this change of the angular anisotropy can be traced to two dynamical origins: the decrease of the overall anisotropy of the main dynamical feature and the appearance of an isotropic slow component.

The most interesting change of the dynamics in this wavelength range is the appearance of a slow peak starting from 225 nm. As the photolysis energy increases, this peak seems to become more important. From the translational energy dependent anisotropy parameter at the photolysis wavelength of 213 nm (Fig.5(b)), the anisotropy parameter is almost zero in the low translational energy region. This is obviously due to the fact that the angular distribution of the slow component is isotropic. At higher translational energy, the anisotropy parameter is about $-0.4$ at this photolysis wavelength, which is also significantly smaller than the value in the low photolysis energy region (230-248 nm). This supports our argument that the overall decrease of
The anisotropy parameter is partly due to the anisotropy decrease of the main dynamics feature at higher translational energies.

The appearance of the slow feature in this wavelength regime turns out to be dynamically quite interesting. Recent theoretical studies show that the energy of appearance is coincident with the threshold of the ring closure pathway of the HN$_3$ molecule, suggesting that the slow feature corresponds to the cyclic N$_3$ product [27]. This is also consistent with the fact that the product angular distribution for this feature is isotropic.

C. Photodissociation dynamics from 208 nm to 190 nm

TOF spectra of the H-atom product were also measured at the even shorter photolysis wavelengths. Four TOF spectra at the photolysis wavelengths of 208, 199, 193.2, and 190 nm with the detector at both the perpendicular and parallel directions to the laser polarization are shown in Fig.9. The photolysis laser polarization was also fixed to be perpendicular to the molecular beam direction for convenience in this wavelength region. The product translational energy distributions converted from these TOF are shown in Fig.10. From these translational energy distributions, translational energy product anisotropy parameters can also be determined. From Fig.6, the overall product anisotropy parameter increases starting from 208 nm. Figure 5(c) shows the typical translational energy dependent anisotropy parameters at the photolysis wavelength of 193.3 nm. At the low translational energy region, the anisotropy parameter remains to be zero as in 217 nm photolysis. The increase of the overall anisotropy is due to the increase of the anisotropy of the fast component. In Fig.5(c), the maximum anisotropy parameter at higher translational energies is about $-0.6$, noticeably larger than that at the 217 nm photolysis. This can be likely due to that fact that different electronic states $B$ and $C$ are becoming accessible at shorter wavelengths.

In comparison with previous photodissociation study at 193.3 nm by Zhang et al. [16], noticeable difference was found in both the transitional energy distribution and the energy dependent angular anisotropy between the result of this work and that by Zhang et al. During the experiment in this work, we found the HN$_3$ photodissociation around 193.3 nm can easily be complicated by multiphoton processes. Therefore, we tried to lower the photolysis laser power as much as possible to reduce the multiphoton effect to minimum in the experiment.
IV. CONCLUSION

Ultraviolet photodissociation of HN₃ has been investigated in the photolysis wavelength range of 188 nm to 264.3 nm using the H-atom Rydberg tagging time-of-flight method. Time-of-flight spectra of the H atom product were measured at the detection directions both perpendicular and parallel to the photolysis laser polarization direction. Total product translational energy distributions as well as product angular anisotropy parameters were determined for the H+N₃ channel at total 12 different photolysis wavelengths. The vibrational state distributions of the linear N₃ product has also been derived at the photolysis wavelengths from 230 nm to 254 nm. Above photolysis wavelength 225 nm, HN₃ predominantly dissociate through the repulsive state and the linear N₃ product vibrational state distributions have been determined for this region. Below 225 nm, a new slow channel starts to appear in addition to the existing direct channel. The appearance energy of this channel is coincident with the theoretical energy threshold of the ring closure pathway of the HN₃ molecule, suggesting that a cyclic N₃ formation is likely present. At even higher photolysis energies, the change of dissociation dynamics is clearly noticeable. This is likely due to the fact that the different electronic excited states are excited at higher photolysis energies.

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