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Rate coefficients for reactions of ethynyl radical (C₂H) with HCN and CH₃CN: Implications for the formation of complex nitriles on Titan

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Abstract. Rate coefficients for the reactions of C₂H + HCN → products and C₂H + CH₃CN → products have been measured over the temperature range 262–360 K. These experiments represent an ongoing effort to accurately measure reaction rate coefficients of the ethynyl radical, C₂H, relevant to planetary atmospheres such as those of Jupiter and Saturn and its satellite Titan. Laser photolysis of C₂H₂ is used to produce C₂H, and transient infrared laser absorption is employed to measure the decay of C₂H to obtain the subsequent reaction rates in a transverse flow cell. Rate constants for the reaction C₂H + HCN → products are found to increase significantly with increasing temperature and are measured to be (3.9–6.2) × 10⁻¹³ cm³ molecules⁻¹ s⁻¹ over the temperature range of 297–360 K. The rate constants for the reaction C₂H + CH₃CN → products are also found to increase substantially with increasing temperature and are measured to be (1.0–2.1) × 10⁻¹² cm³ molecules⁻¹ s⁻¹ over the temperature range of 262–360 K. For the reaction C₂H + HCN → products, ab initio calculations of transition state structures are used to infer that the major products form via an addition/elimination pathway. The measured rate constants for the reaction of C₂H + HCN → products are significantly smaller than values currently employed in photochemical models of Titan, which will affect the HC₃N distribution.

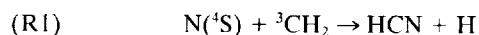
1. Introduction

Over the last 2 years there have been two new photochemical models of Titan's atmosphere [Lara *et al.*, 1996; Toubanc *et al.*, 1995]. These models differ from the first post-Voyager model [Yung *et al.*, 1984; Yung, 1987] primarily in the updated methane photodissociation scheme [Mordaunt *et al.*, 1993], better characterizations of the eddy diffusion coefficient, and updated reaction rates. Gladstone *et al.* [1996] also presented a photochemical model for the upper atmosphere of Jupiter based on similar reaction sequences.

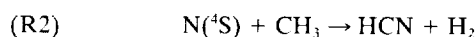
With the upcoming Cassini/Huygens mission to Saturn and its largest satellite Titan, there is renewed interest in understanding the photochemical processes that take place in the atmosphere of Titan. This interest generally arises from the presence of a dense N₂-CH₄ atmosphere which is thought to possess many similarities with the prebiotic chemistry of the Earth [Raulin *et al.*, 1994]. Efforts to unravel the processes that lead to the formation of life are not new, with attempts to simulate Earth's primitive atmosphere originating over 40 years ago [Miller, 1953]. Numerous experiments have been carried out to simulate the synthesis of organic molecules in Titan's atmosphere, including UV radiation, high-energy particles (H⁺, e, γ), and electric discharges. These laboratory simulations of Titan's atmosphere produce a remarkable mixture of C₂ through C₆ compounds as well as various nitriles [Cabane and Chassefière, 1995].

Hydrogen cyanide (HCN), cyanoacetylene (HC₃N), and cyanogen (C₂N₂) have been identified as components of Titan's

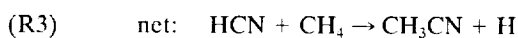
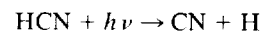
atmosphere with the Voyager 1 infrared spectrometer (IRIS) [Coustenis, 1990; Coustenis *et al.*, 1991], while ethanenitrile (CH₃CN), commonly referred to as acetonitrile, methyl cyanide, or cyanomethane, has been detected via ground-based millimeter spectroscopy [Bézar *et al.*, 1993]. The models of Lara *et al.* [1996], Toubanc *et al.* [1995], and Yung *et al.* [1987] and Yung [1984] postulate that HCN is formed from the reaction of ground state nitrogen atoms when they react with ground state methylene:



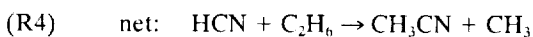
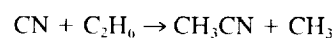
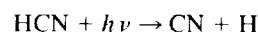
Yung *et al.* also included the reaction of ground state nitrogen with methyl radicals.



Toubanc *et al.* [1995] and Lara *et al.* [1996] proposed the following mechanism for the formation of CH₃CN:



Lara *et al.* also included a second photochemical process involving ethane:



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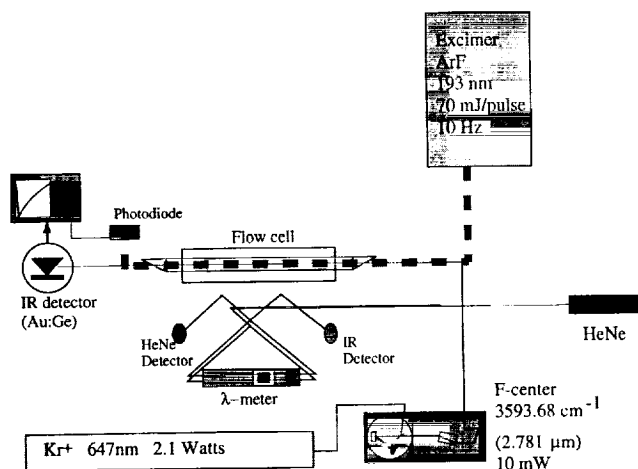
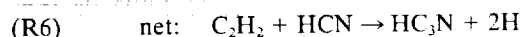
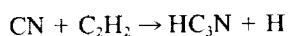
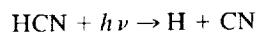
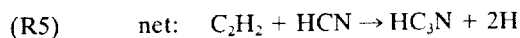
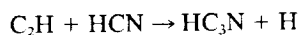
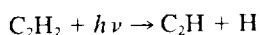


Figure 1. Schematic of experimental setup.

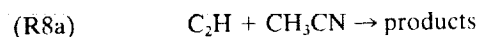
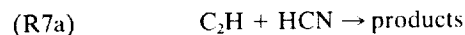
The main loss processes for CH₃CN are considered to be photolysis and vapor condensation.

These compounds, as well as acetylene and the corresponding radicals, are the main chemical species thought to be involved in the production of complex nitriles. Each of the above models for Titan include the following two schemes for the production of cyanoacetylene, HC₃N:

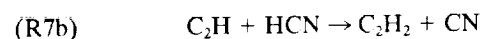


The competing, exothermic H atom abstraction reaction, C₂H + HCN → C₂H₂ + CN, was not considered.

The current research is focused on the reactivity of the ethynyl radical, C₂H, with HCN and CH₃CN:



There are no experimental data to define the products of these reactions. Both hydrogen abstraction reactions and addition/elimination processes are thermodynamically possible:



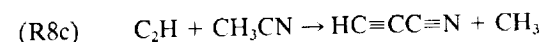
$$\Delta H_{rxn} = -38 \text{ kJ mol}^{-1}$$



$$\Delta H_{rxn} = -126 \text{ kJ mol}^{-1}$$



$$\Delta H_{rxn} = -180 \text{ kJ mol}^{-1}$$



$$\Delta H_{rxn} = -150 \text{ kJ mol}^{-1}$$

(Values of ΔH_{rxn} are calculated on the basis of available ΔH_f^0 values from *Weast* [1990] except for cyanoacetylene, which is from *Okabe and Dibeler* [1973].)

The measurements presented here for the reaction rates of C₂H with the nitrogen containing species, HCN and CH₃CN, are the first direct measurements of the total rate coefficients for these reactions. Since acetylene and polyynes are some of the primary photochemically active species in the atmosphere of Titan, the coupling of the resulting C₂H radicals with the potentially reactive nitrogen compounds is important. In this paper, rate constants for the reactions of C₂H with HCN and CH₃CN are obtained; possible product pathways and the impact of the results on photochemical models are discussed.

2. Experiment

The experimental technique used in the determination of low-temperature rate coefficients of the ethynyl radical has been described in detail [Pedersen *et al.*, 1993; Opansky and Leone, 1996], and only a brief overview is presented here. A schematic of the experimental arrangement is shown in Figure 1. The ethynyl radicals are produced by photolysis of acetylene in a flow cell using a pulsed excimer laser operating at 193 nm (70 mJ/pulse at 10 Hz). The transient decay of the C₂H concentration is probed by a single-mode infrared F-center laser operating at 3593.68 cm⁻¹, which is used to monitor the absorption of the Q₁₁(9) line of the A²Π – X²Σ⁺ (000) transition. The longitudinal mode structure of the F-center laser is monitored with a scanning Fabry-Perot spectrum analyzer, and a traveling wave meter is used for measuring the wavelength [Hall and Lee, 1976].

The kinetic experiments are performed in a 1 m long, 2.5 cm diameter, variable temperature flow cell. The photolysis and probe lasers are overlapped over the full length of the cell, and a three-times multipass of the probe beam is employed to increase the absorption path length. Gases are introduced into and removed from the cell via a series of transverse inlets and pump-outs, which allows rapid removal of the photolysis products from the probe volume. The cell was cooled by the circulation of pentane or isobutane through an outer jacket after having passed through a copper coil submerged in an ethanol/liquid nitrogen slush bath. Temperatures of ≈155 K are easily obtained with this method. Due to the slowness of the C₂H + HCN reaction and the limited vapor pressure of CH₃CN, measurements were restricted to low temperatures of 297 K for HCN and 262 K for CH₃CN. Measurements above room temperature to ≈360 K are made by circulating water through the outer jacket while the copper coils are placed in a water reservoir in which the temperature is controlled by an immersion heater. The temperature in the flow cell is measured by three K-type thermocouples positioned along the length of the cell, and partial pressures of each gas are calculated from the measured flow rates using calibrated mass flow meters and the total pressure. Typical number densities for C₂H₂ are (2–6) × 10¹⁴ cm⁻³; number densities for HCN and CH₃CN are (0.5–3) × 10¹⁶ cm⁻³ and (2–8) × 10¹⁶, respectively, with a buffer gas of helium in large excess. Using an absorption cross section for C₂H₂ at 193 nm of 1.35 × 10⁻¹⁹ cm² and a quantum yield of 0.26 [Satyapal and Bersohn, 1991], the upper limit for [C₂H] is estimated to be 6.5 × 10¹¹ cm⁻³. The total pressure for typical data acquisition on the CH₃CN reaction is between 3.7 and 4.1 kPa (28–31 torr); pressure dependence measurements are conducted over the range of 1.3–12 kPa (10–90 torr). Reaction

rates were independent over this pressure range. The total pressure for data acquisition on the HCN reaction is between 3.7 and 12 kPa (28–90 torr), and the reaction was assumed to be independent of pressure over this range. This large pressure range was required because of the use of a 1% HCN mixture in He; therefore the number density of He is directly related to the possible HCN concentrations obtained. All gases flow through a mixing cell before entering the low-temperature cell. He, 99.99%, C₂H₂, 99.6%, and HCN, 1.0% in helium, are obtained from commercial sources, and most gases are used as received, except acetylene, from which acetone was removed by an activated charcoal filter. Commercially available liquid CH₃CN, 99.9%, was vacuum distilled and degassed.

The probe beam is directed onto a 50 MHz liquid N₂ cooled Ge: Au infrared detector with a sensitive area of 20 mm². The transient signals are amplified and co-added using a digital oscilloscope to improve the signal-to-noise ratio. About 1500–3000 transient absorption signals are averaged before the data are transferred to a personal computer, where the data are stored and analyzed.

3. Results

Rate coefficients are measured under pseudo-first-order conditions where [HCN], [CH₃CN], and [C₂H₂] are ≫ [C₂H]. The rate of change for [C₂H] can be expressed as

$$d[\text{C}_2\text{H}]/dt = -[\text{C}_2\text{H}](k_{\text{RCN}}[\text{RCN}] + k_{\text{acetylene}}[\text{C}_2\text{H}_2]) \quad (1)$$

where R represents either H or CH₃. After integration of (1),

$$[\text{C}_2\text{H}] = [\text{C}_2\text{H}]_0 \exp[-k_{\text{obs}}(t)] \quad (2)$$

where

$$k_{\text{obs}} = k_{\text{RCN}}[\text{RCN}] + k_{\text{acetylene}}[\text{C}_2\text{H}_2] \quad (3)$$

$$k_{\text{obs}} - k_{\text{acetylene}}[\text{C}_2\text{H}_2] = k_{\text{RCN}}[\text{RCN}] = k'_{\text{RCN}} \quad (4)$$

The observed decay rates of C₂H, k_{obs} , are obtained by fitting the transient signal to the following equation:

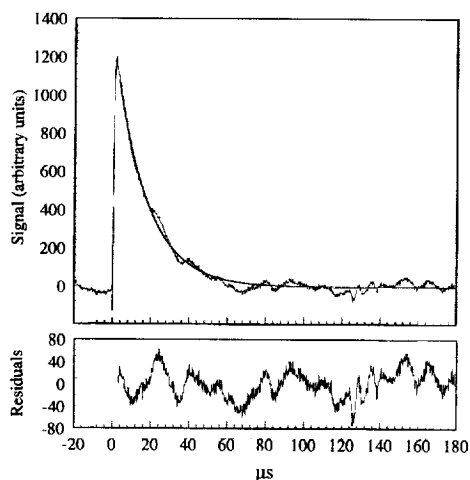


Figure 2. (upper trace) Typical C₂H transient absorption signal at $T = 360$ K, total pressure 12.3 kPa, $[\text{C}_2\text{H}_2] = 2.8 \times 10^{14}$ molecules cm^{-3} , $[\text{HCN}] = 2.5 \times 10^{16}$ molecules cm^{-3} , and $k_{\text{obs}} = 0.059 \pm 0.001 \mu\text{s}^{-1}$ (uncertainty represents 95% confidence limit). (lower trace) Residuals to fitted experimental decay.

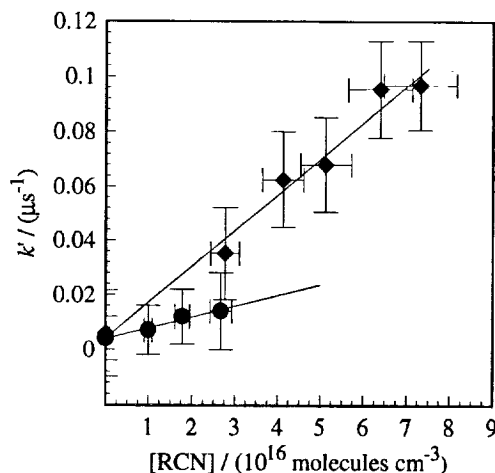


Figure 3. Plot of k'_{RCN} versus $[\text{RCN}]$: diamonds, CH₃CN at $T = 296$ K and $k_{\text{CH}_3\text{CN}} = (1.3 \pm 0.2) \times 10^{-12}$ cm^3 molecules⁻¹ s⁻¹; circles, HCN at $T = 297$ K and $k_{\text{HCN}} = (3.9 \pm 1.2) \times 10^{-13}$ cm^3 molecules⁻¹ s⁻¹.

$$y = A \exp[-k_{\text{obs}}(t)] + \text{const} \quad (5)$$

An example of a typical C₂H experimental decay and corresponding fit for the reaction with HCN is shown in Figure 2. The values of k_{obs} are corrected for the C₂H + C₂H₂ contribution by subtraction in (4) using the most recent measurements for $k_{\text{acetylene}}$ [Opansky and Leone, 1996]. The contribution of $k_{\text{acetylene}}$ to k_{obs} is between 35 and 70% for the determination of k_{HCN} and less than 40% for the determination of $k_{\text{CH}_3\text{CN}}$. The resulting k'_{RCN} values are plotted as a function of concentration and fit using a linear least squares routine. The resultant slope is equal to k_{RCN} ; this is shown in Figure 3 for HCN and CH₃CN. The uncertainty in k_{RCN} is determined by computing the standard deviation of the mean for each of the plots of k_{RCN} . The reported errors are $\pm 2\sigma$. The experimental results for C₂H + HCN and CH₃CN are summarized in Table 1, and the corresponding Arrhenius plots are shown in Figure 4. The data can be expressed using the Arrhenius forms $k_{\text{RCN}} = A \exp[(E_a)/(RT)]$ as $k_{\text{HCN}} = (5.3 \pm 2.2) \times 10^{-12} \exp[(-6.4 \pm 1.2 \text{ kJ/mol})/(RT)]$ and $k_{\text{CH}_3\text{CN}} = (1.8 \pm 1.2) \times 10^{-11} \exp[(-6.4 \pm 2.0 \text{ kJ/mol})/(RT)]$. The relatively large uncertainty of the Arrhenius parameters is due to the narrow temperature range of the experiments. The observed transient signals can be fit to first-order exponential decays. The standard deviation of each rate constant at a given temperature is taken as the overall experimental uncertainty of each data point (20–30%). The most likely source of systematic error is a reduction in the actual number density of the reactant species due to condensation. The ex-

Table 1. Summary of Rate Constants for C₂H + RCN

Temperatures, K	k_{HCN}	Temperature, K	$k_{\text{CH}_3\text{CN}}$
297	$(3.9 \pm 1.2) \times 10^{-13}$	262	$(1.0 \pm 0.2) \times 10^{-12}$
330	$(4.9 \pm 1.3) \times 10^{-13}$	296	$(1.3 \pm 0.2) \times 10^{-12}$
360	$(6.2 \pm 1.8) \times 10^{-13}$	296	$(1.2 \pm 0.2) \times 10^{-12}$
		360	$(2.2 \pm 0.3) \times 10^{-12}$

Here $k_{\text{HCN}} = (5.3 \pm 2.2) \times 10^{-12} \exp[(-6.4 \pm 1.2 \text{ kJ/mol})/(RT)]$; $k_{\text{CH}_3\text{CN}} = (1.8 \pm 1.2) \times 10^{-11} \exp[(-6.4 \pm 2.0 \text{ kJ/mol})/(RT)]$. Values are in units of cm^3 molecules⁻¹ s⁻¹.

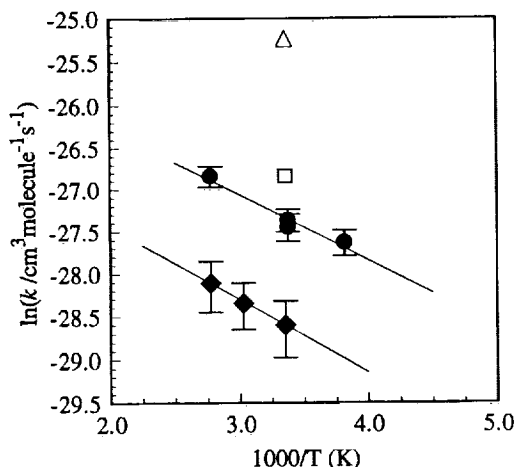


Figure 4. Arrhenius plot: Diamonds, C₂H + HCN; solid line, $k_{\text{HCN}} = 5.3 \times 10^{-12} \exp [(-6.4 \text{ kJ/mol})/(RT)] \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$; circles, C₂H + CH₃CN; solid line, $k_{\text{CH}_3\text{CN}} = 1.8 \times 10^{-11} \exp [(-6.4 \text{ kJ/mol})/(RT)] \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$; open squares, rate constant for the reaction C₂H + HCN → HC₃N + H used in the photochemical models of Yung *et al.* [1984], Yung [1987], and Toubanc *et al.* [1995]; and open triangle shows rate constant for the reaction C₂H + HCN → HC₃N + H used in the photochemical model of Lara *et al.* [1996].

periments involving HCN were all performed at temperatures ≥ 297 K, where HCN does not condense; however, in the case of CH₃CN, attempts to measure rate constants below 262 K resulted in erratic pressure measurements, attributed to condensation of CH₃CN in the flow cell.

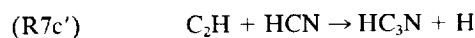
Under the current experimental conditions, the minimum C₂H₂ density necessary for detection of the C₂H radical is approximately 1×10^{11} molecules cm⁻³. The rate constant for the reaction of C₂H with the precursor C₂H₂ is rapid, 1.36×10^{-10} cm³ molecule⁻¹ s⁻¹ at 298 K [Pederson *et al.*, 1993; Opansky and Leone, 1996]. The combination of this extremely fast reaction and the high density of HCN needed to make rate coefficient measurements presents a serious limitation in the current experiment. Thus we are able to measure the HCN reaction only at temperatures of 297 K and above. An attempt to measure the rate constant at 200 K was made to determine whether the rate might show non-Arrhenius behavior and increase again at even lower temperatures; however, there was no evidence that the rate increases at this temperature, and only the experimental decay from the reaction of C₂H with the precursor molecule acetylene was observable. Use of a pure sample of HCN would allow for higher densities of the reactant and thus increase the observed first-order decay rate; however, this would also lead to larger concentrations of CN radicals from the resulting HCN photolysis (under the current experimental conditions, the CN radical concentration is estimated to be of the same order as C₂H). Another solution would be to improve the sensitivity of the experimental arrangement, by implementing a two-detector signal subtraction scheme to reduce noise from amplitude fluctuations in the laser. Nevertheless, limitations due to the HCN vapor pressure would also make lower temperature measurements difficult. This vapor pressure limitation is observed for CH₃CN. Although the lowest possible experimental temperature is approximately 150 K, rate constants for CH₃CN could only be determined above 262

K. Measurements at temperatures below 262 K were not possible because of condensation of the CH₃CN vapor.

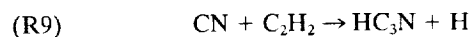
4. Discussion

The experimental results for the reactions of C₂H + HCN and C₂H + CH₃CN show positive temperature dependencies, indicating barriers to reaction, as well as small rate coefficients.

There has been one previous study that examined only the overall product photochemistry of acetylene and hydrogen cyanide mixtures [Becker and Hong, 1983]. In those experiments, photolysis of C₂H₂ and HCN mixtures were exposed to 185 nm light from a low-pressure mercury lamp (10^7 photons/s) for 2 min. Gaseous products were then analyzed by gas chromatography/mass spectroscopy. Photolysis products for 5:1 mixtures of HCN/C₂H₂ were similar to those of C₂H₂ (where C₄H₂, diacetylene, was the major product) except for the formation of a significant amount of acrylonitrile, C₂H₃CN. Trace amounts of HC₃N were reported. Under these conditions, only 10% of the light is absorbed by HCN. At higher HCN/C₂H₂ ratios, where 40–60% of the light is absorbed by HCN, HC₃N becomes a major product. As discussed above, the formation of HC₃N is believed to be generated by the possible reactions



$$\Delta H_{\text{rxn}} = -126 \text{ kJ mol}^{-1}$$



$$\Delta H_{\text{rxn}} = -88 \text{ kJ mol}^{-1}$$

Becker and Hong concluded that reaction (R9) was the major process for the formation of HC₃N in their photochemical studies, even though both reactions are exothermic.

The temperature dependence of the kinetic rate coefficients can provide information on the possible mechanisms of the chemical reactions. Although data have been obtained at only three temperatures, the rate constant for C₂H + HCN is decreasing substantially with decreasing temperature. Attempts to measure rate constants at temperatures below 297 K were not possible because of the slow rate of the reaction. A positive temperature dependence and large activation energy are typical of either hydrogen abstraction reactions or addition/elimination reactions that proceed over a barrier in the reaction coordinate. An Arrhenius plot that has little or no temperature dependence or a negative temperature dependence is often associated with an addition/elimination reaction.

In order to better understand the possible pathways for the reaction C₂H + HCN → products, ab initio calculations have been carried out on the transition states corresponding to both the addition/elimination process (products = HC₃N + H) and the hydrogen abstraction mechanism (products = C₂H₂ + CN). The calculations are carried out with a commercial software package using the 6-31 g (d,p) basis set [Frisch *et al.*, 1995]. Geometries for both transition states and individual molecules are calculated using full Møller-Plesset second-order perturbation theory (MP2). Transition state structures are confirmed in a corresponding vibrational frequency calculation [Foresman and Frisch, 1996]. Final energies are calculated using the quadratic configuration interaction with single and double excitation substitutions (QCISD). The results of these calculations are shown in Figure 5. Classic barriers are determined without corrections for zero-point or thermal energies. The results show that the transition state corresponding

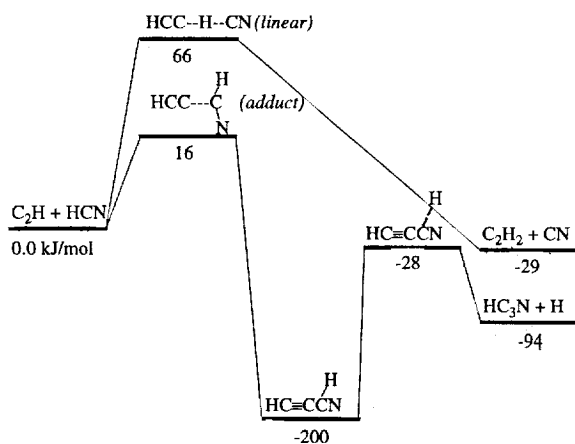
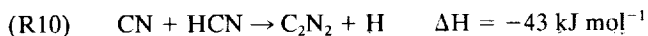


Figure 5. Energy diagram showing classic barrier heights for the reaction $C_2H + HCN \rightarrow$ products. Calculations carried out using QCISD/6-31g(d,p)//MP2(full)/6-31g(d,p) method.

to the hydrogen abstraction mechanisms lies approximately 43 kJ mol^{-1} (10 kcal mol^{-1}) higher in energy than the transition state for the addition/elimination mechanism, which suggests that the addition/elimination mechanism is the dominant reaction pathway. Although the calculated classic barrier of 15.9 kJ mol^{-1} (3.8 kcal mol^{-1}) overestimates the experimental value of 6.4 kJ mol^{-1} (1.5 kcal mol^{-1}), the presence of a barrier along the reaction coordinate agrees qualitatively with the observed positive temperature dependence.

The measurements for the reaction of C_2H with CH_3CN give rate constants that are very close to those measured previously for methane, CH_4 ($k_{\text{methane}} = 1.2 \times 10^{-11} \exp [(-4.1 \text{ kJ/mol})/(RT)]$; $154 < T < 359$). Over the observed temperature range, the reaction has a positive temperature dependence and large activation energy. The slower reaction rate for CH_3CN (compared to CH_4) could represent a steric factor due to the presence of the CN group of the CH_3CN ; however, based on the experimental measurements and calculations for the $C_2H + HCN$ reaction, an addition/elimination mechanism is likely to contribute to the observed rate constant for C_2H with CH_3CN .

Since this is the first experimental work to directly measure the rate coefficients for reactions (R7a), it is useful to make the comparison with the analogous reactions involving the CN radical, which is isoelectronic with C_2H :



Yang *et al.* [1992] measured rate coefficients for reaction (R10) over the temperature range of 300–740 K. They report room temperature rate constants of $\approx 2.6 \times 10^{-14} \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$ and fit their results to the modified Arrhenius equation

$k_{R10} = 2.5 \times 10^{-17} T^{1.71} \exp(-770/T) \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$. In those experiments, only the disappearance of CN was monitored, and no information was obtained on the product channels. The result corresponds to the present work on $C_2H + HCN$, in that the rate constant for the addition/elimination reaction is relatively small and the reaction shows a strong positive temperature dependence. Yang *et al.* [1992] calculated a reaction barrier of 13.8 kJ mol^{-1} (3.3 kcal mol^{-1}) for the reaction $CN + HCN \rightarrow H(CN)_2 \rightarrow H + C_2N_2$.

5. Implications for Formation of Cyanoacetylene on Titan

We first focus on the implications of this work for the chemical formation of cyanoacetylene, HC_3N . Photochemical models of Titan have used various rate constants for reactions (R7c), $C_2H + HCN \rightarrow HC_3N + H$ and (R9), $CN + C_2H_2 \rightarrow HC_3N + H$, which are summarized in Table 2. The rate coefficients used in the models for (R7c), also shown in Figure 4, can be compared to the experimental results obtained here. The results show that the rate constant used in the most recent photochemical model [Lara *et al.*, 1996] is too fast by almost a factor of 30 when compared to the room temperature values obtained here. The two earlier models [Yung *et al.*, 1984; Yung, 1987; Toubanc *et al.*, 1995] use a value which is only a factor of 5 faster than the measured room temperature results. If we make the assumption that the reaction follows the measured Arrhenius behavior, the expected rate constant would be $\approx 1 \times 10^{-13}$ at 200 K and $\approx 5 \times 10^{-14}$ at 170 K (the latter temperature being close to the value of the lower stratosphere on Titan) and the discrepancy only increases.

The photochemical models of Titan make comparisons of their vertical distributions of cyanoacetylene with available Voyager/IRIS data. Lara *et al.* [1996] pointed out a discrepancy between the mixing ratio of HC_3N determined by Voyager/IRIS data and the value in their model profiles. The results showed that the concentration and column density depended strongly on the eddy diffusion coefficient. Using the eddy diffusion coefficient proposed in their model resulted in vertical profiles for the HC_3N mixing ratio that were too high by nearly 2 orders of magnitude when compared with the available Voyager/IRIS data. Lara *et al.* noted that the most important reactions in their modeling of HC_3N production are reactions (R7c) and (R9). As mentioned above, the rate constant for (R7c) used in their photochemical model is nearly a factor of 30 faster than the value measured in the current experiments. That value of the rate constant was recommended by Monks *et al.* [1993] assuming analogous behavior with the addition/elimination reaction $C_2H + C_2H_2$ (C_2H_4) $\rightarrow C_4H_2$ (C_4H_4) + H. The overprediction of HC_3N may be further aggravated by the rate constant used for reaction (R9). Using the non-Arrhenius expression given by Lara *et al.* [1996], the rate cal-

Table 2. Comparison of Experimental Rate Constants for Reactions Leading to HC_3N With Values Used in Recent Photochemical Models of Titan

Reaction	Experiment	Yung <i>et al.</i> [1984]	Yung [1987]	Toubanc <i>et al.</i> [1995]	Lara <i>et al.</i> [1996]
$C_2H + HCN \rightarrow HC_3N + H$	$5.3 \times 10^{-12} e^{(-770/T)}$ ^a	2.2×10^{-12}	...	2.2×10^{-12}	1.1×10^{-11}
$CN + C_2H_2 \rightarrow HC_3N + H$	$\approx 4 \times 10^{-10}$ ^b	5×10^{-11}	2.3×10^{-11}	$3.49 \times 10^{-11} e^{(571/T)}$	$5.67 \times 10^{-9} T^{0.55} e^{(-4/T)}$

Values are in units of $\text{cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$.

^aCurrent results.

^bFrom Sims *et al.* [1993], $25 < T < 80$ K.

Table 3. Parameters and Production Rates of HC₃N Via C₂H + HCN and CN + C₂H₂ Reaction Pathways

Parameter	<i>Yung et al.</i> [1984]	<i>Toublanc et al.</i> [1995]	<i>Lara et al.</i> [1996]	
[C ₂ H] at 750 km	≈3 × 10 ³ molecules cm ⁻³			
[CN] at 750 km	≈1 × 10 ² molecules cm ⁻³			
Mixing ratio HCN	10 ⁻⁴	10 ⁻⁴	2 × 10 ⁻⁵ at 400 km ^a	
Mixing ratio C ₂ H ₂	7 × 10 ⁻⁴	1.5 × 10 ⁻³	8 × 10 ⁻⁴	
k(C ₂ H + HCN) ^b	2.2 × 10 ⁻¹²	2.2 × 10 ⁻¹²	1.1 × 10 ⁻¹¹	
k(CN + C ₂ H ₂)	5 × 10 ⁻¹¹	1 × 10 ⁻¹¹ (T = 170 K)	9 × 10 ⁻⁸ (T = 170 K)	
n(750 km)	5 × 10 ¹¹ molecules cm ⁻³			
Production Rates of HC ₃ N, molecules cm ⁻³ s ⁻¹				
Reaction Pathway	This work	<i>Yung et al.</i> [1984]	<i>Toublanc et al.</i> [1995]	<i>Lara et al.</i> [1996]
C ₂ H + HCN → HC ₃ N + H	0.008 ^c	0.3	0.5	0.3
CN + C ₂ H ₂ → HC ₃ N + H	14 ^d	2	75	3600

^aAssumed to have small variance between 400 and 750 km.

^bAll rate constants have units cm³ molecules⁻¹ s⁻¹.

^cUsing $k = 5 \times 10^{-14}$ (from Arrhenius expression at 170 K based on this work).

^dUsing $k = 4 \times 10^{-10}$ [from *Sims et al.*, 1993].

culated at 170 K would be 9.3×10^{-8} cm³ molecules⁻¹ s⁻¹. This rate is nearly 2 orders-of-magnitude faster than the gas-kinetic rate. In addition, *Sims et al.* [1993] recently reported the rate coefficient for reaction (R9) down to 25 K. At temperatures below 100 K the rate constant is $\approx 4 \times 10^{-10}$ cm³ molecules⁻¹ s⁻¹, very close to the gas-kinetic rate. Between 100 and 200 K the reaction rate is still very fast, $2\text{--}3 \times 10^{-10}$ cm³ molecules⁻¹ s⁻¹. The large difference in the rate used in the model may be responsible for the source of discrepancy in the computed vertical distributions of HC₃N.

The models of *Yung et al.* [1984; *Yung*, 1987] and *Toublanc et al.* [1995] used the same, estimated value for the rate constant of reaction (R7c), C₂H + HCN → HC₃N + H; however, their use of significantly different values of the rate constant for reaction (R9), CN + C₂H₂ → HC₃N + H, make direct comparison between the models difficult. Other factors in addition to the difference in rate constants (e.g., the eddy diffusion coefficient, reactivity of HC₃N, photolysis rate) contribute to the observed differences in the vertical profiles of HC₃N. *Coustonis* [1990] sets an upper limit for the equatorial mixing ratio for HC₃N at 105 km of $\leq 1.5 \times 10^{-9}$. At the time of the *Yung et al.* [1984] models, only crude estimates were available for the concentration of this species, and their column-averaged abundances for HC₃N were in good agreement with the available data [*Kunde et al.*, 1981]. Examination of the *Yung et al.* [1984] HC₃N vertical profile indicates that the mixing ratio is approximately 10⁻⁷ at 100 km. However, *Toublanc et al.* [1995] indicate that their mixing ratio of HC₃N at 105 km is $\leq 10^{-12}$, although the stratospheric abundances (450–850 km) of *Toublanc et al.* [1995] appear to be within a factor of 10 of those of *Yung et al.* [1984]. The nitrile photochemistry of *Yung et al.* [1984] was revised [*Yung*, 1987] mainly to include the updated rate constant of the reaction CN + HCN → C₂N₂ + H and new chemical schemes for the formation of cyanogen, C₂N₂. However, the overall vertical distributions for the nitrile compounds included in the model changed little from the original work due to the inclusion of additional pathways for the formation of C₂N₂.

An evaluation can be carried out to quantify the results presented here in terms of the production rate of HC₃N. The reaction rates for C₂H + HCN and CN + C₂H₂ were calculated for concentrations of molecular species at 750 km. (This

altitude was chosen because the mixing ratio of C₂H was at a maximum in the *Yung et al.* [1984] paper.) Concentrations for the radical species C₂H and CN are from *Yung et al.* [1984]. This is the only reference that reports the vertical profiles of these radical species. Concentrations of the molecular species are determined by multiplying the reported mixing ratios (mole fractions) by the total number density at 750 km [*Yung et al.*, 1984]. The data required for this comparison as well as the production rates are shown in Table 3. The models assume that each reaction has only one product channel. This allows one to set the reaction rate for each process equal to the production rate of HC₃N for the particular reaction pathway. It is interesting to note the similarities of the three models for the production of HC₃N via the C₂H + HCN pathway; however, a major difference occurs between the model values and the value obtained using the new rate constant of 5×10^{-14} cm³ molecules⁻¹ s⁻¹ (170 K) extrapolated from the Arrhenius expression determined in the present study. Similar estimations were made for the CN + C₂H₂ reaction pathway. It is clear that the CN + C₂H₂ reaction dominates in the production of HC₃N.

It is important to note that while the current paper deals with differences in the rate constants obtained experimentally and those used in photochemical models, other factors also influence the calculated vertical distributions of HC₃N. Processes such as diffusion, mixing as well as loss processes associated with condensation, and chemical reactivity are also important in predicting the concentration of minor atmospheric constituents; however, the use of accurate reaction rates is essential if a complete understanding of the photochemical processes is to be achieved.

Finally, current photochemical models include CH₃CN but limit the loss processes to photolysis and condensation. Since the concentration of CH₃CN on Titan is low and the rate constant presented here is relatively small, the inclusion of the reactions of C₂H with CH₃CN would most likely have little effect on the predicted vertical profiles; however, the reaction C₂H + CH₃CN → HC₃N + CH₃ does represent a possible additional pathway for the formation of cyanoacetylene, HC₃N.

The variations between the three models demonstrate the need for further refinement in both experimental and theoret-

ical work. The use of updated rate coefficients such as those presented here and by other research groups, as well as a prudent analysis of the reaction pathways should allow future models to better estimate atmospheric abundances of major and minor constituents on Titan. The upcoming Cassini/Huygens mission will greatly enhance our current knowledge of Titan's atmosphere.

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