Astronomical detection of the cyanobutadiynyl radical C$_5$N

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Abstract. We report the detection of the elusive carbon-chain radical C$_5$N in the dark cloud TMC1 and its tentative detection in the circumstellar envelope IRC+10216. C$_5$N appears to be two orders of magnitude less abundant than the related molecule HC$_5$N and much less abundant than expected from current gas phase chemistry models. In comparison the HC$_3$N to C$_5$N abundance ratio is of the order of 10, in reasonable agreement with model predictions.

We have also detected in IRC+10216 two lines arising from the C$_3$H radical in its excited $\nu_4=1$ state.

Key words: Molecular data – circumstellar matter – ISM: molecules – Radio lines: stars

1. Introduction

That long carbon chain radicals could be abundant and play a large role in interstellar chemistry was first recognized with the discoveries of C$_3$N in the circumstellar envelope IRC+10216 (Guélin & Thaddeus 1977) and in the dark cloud TMC1 (Friberg et al. 1980). To date seven acetylenic chain radicals, C$_n$H, $n=2–8$, and five cyanopolyne molecules, HC$_{2n}$CN, $n=1–5$, are identified in TMC1 and/or IRC+10216 (Bell et al. 1997, Guélin et al. 1997). Surprisingly, no cyanopolyne radicals heavier than C$_3$N were so far detected, despite model predictions that at least C$_5$N should be abundant (Herbst et al. 1994).

The non-detection of C$_5$N was first blamed on a small dipole moment and on the lack of spectroscopic data. The dipole moment $\mu_0$ and the rotation constant $B_0$ of the C-chains radicals depend critically on the nature of their electronic ground state. The two lowest states, $^2\Pi$ and $^2\Sigma$, are close in energy (Pauzat et al. 1991). Pauzat and co-workers predicted from unrestricted Hartree-Fock calculations that the $^2\Pi$ state of C$_5$N lay below the $^2\Sigma$ state and that $\mu_0$ was very small. The line strengths scaling with $\mu_0^2$, it was no wonder that C$_5$N escaped detection.

Things changed when Botschwina (1996) showed from more elaborate coupled cluster calculations that the C$_5$N ground state was in fact $^2\Sigma$ and that $\mu_0$ was as large as 3.385 D, even larger than the dipole moment of C$_3$N. It then became clear that C$_5$N could be detected at least in the laboratory. Kasai et al. (1997) succeeded to synthesize this species, to measure its microwave spectrum, and to derive its rotational, fine and hyperfine constants, making at last possible a sensitive search for C$_5$N in space. In this Letter, we report the astronomical detection of this radical.

2. Observations and results

The long carbon-chain molecules and radicals are nearly thermalized in TMC1 and IRC+10216 with rotation temperatures, $T_{rot}$, in the ranges 6–10 K and 20–50 K, respectively. Since, for C$_3$N, $B_0 B_2^{1/2} = 0.067$ K, the strongest lines in these two sources lie in the $\lambda = 1$ cm and $3–4$ mm atmospheric windows. We thus searched for C$_5$N at these wavelengths, using the Effelsberg 100-m telescope and the Pico Veleta 30-m telescope.

The Effelsberg observations were made in October 1997, January and March 1998. The telescope was equipped with the K-band maser receiver. The weather was mostly clear and the system temperature $T_{sys} \simeq 100$ K. We used the new AK90 autocorrelator split into two 20 Mhz-wide bands of 4048 channels each. One of the bands covered both fine structure components of the $N = 9 \rightarrow 8$ transition of C$_5$N (25.250 GHz, Kasai et al. 1997), the other the $J = 21.5 \rightarrow 20.5$ transition of $^2\Pi_{3/2}$ C$_4$H (25.227 GHz). The local oscillator frequency was switched by $\pm 200$ kHz or $\pm 500$ kHz and the spectra folded accordingly. The average spectrum obtained in TMC1 at the position of the cyanopolyne peak (1950.0, $4^p 38^m 38.6^s, 25^\circ 35^\prime 45^\alpha$), smoothed to 15 kHz (0.17 kms$^{-1}$), has a r.m.s. noise of 3.5 mK (units of $T_{MB}$).

In addition to C$_3$N and C$_4$H, we observed briefly the $J = 10 \rightarrow 9$ line of HC$_5$N (23.96 GHz) and the $N = 2 \rightarrow 1$ lines of C$_3$N (19.79 GHz). All the data were calibrated following the procedure described by Schilke & Walmsley (1991) by observing at every frequency the planetary nebula NGC 7027, whose flux was taken equal to 5.8 Jy. The calibration uncertainty, which results mostly from the atmospheric absorption correction and from beam efficiency...
Detection of $C_5N$

Fig. 1. Spectrum observed with the 100-m Effelsberg telescope toward TMC 1 (cyanopolyyne peak: $4^{h}38^{m}38.6^{s}, 25^\circ35'45''$, 1950.0) and covering the N=9–8 rotational transition of $C_5N$. The N=9–8 line is split into two fine structure components whose frequencies, derived from laboratory measurements, are marked by upward arrows. The hyperfine structure is too small to be resolved.

The spectrum was observed by switching the local oscillator in frequency by $\pm 200$ kHz or $\pm 500$ kHz; the position of the ghosts of the line in the folded spectrum are indicated by downward arrows. The average variation, is estimated to be $< 20\%$. The HC$_5$N and C$_3$N line intensities we measure are consistent with those reported by Tölle et al. (1981) and Guélin et al. (1982).

Figure 1 (see also Table 1) shows the spectrum covering the $C_5N$ transitions. We see two 0.3 kms$^{-1}$-wide spectral lines, each detected at $> 5\sigma$. The lines are separated by $10.71 \pm 0.01$ MHz, which is very close to the value of the spin-rotation constant measured by Kasai et al. (1997), $\gamma = 10.75$ MHz. Their half-power width is comparable to the C$_3N$ linewidth, $0.23 \pm 0.05$ kms$^{-1}$. Their rest frequencies coincide within the small uncertainties with the C$_5N$ N=9–8 transition frequencies calculated by Kasai et al. (1997), if we adopt the source LSR velocity of $5.65 \pm 0.05$ kms$^{-1}$ measured for C$_3$N and C$_3$H (see Guélin et al. 1982). Since there are no other comparable lines in the 20 MHz-wide spectrum we observed, the probability for a chance coincidence is $< 10^{-6}$. We thus conclude that we have detected C$_5N$ in TMC1.

The 30-m observations were made in Nov. 1997 and April 98. The telescope was equipped with two SIS mixer receivers with orthogonal polarizations. The zenith atmospheric opacity was below 0.1 and the system temperature $T_{sys} = 150 - 200$ K. The observations were made by wobbling in azimuth the secondary mirror at a rate of 0.5 Hz and with an amplitude of 90''. We searched for the N=32–31 line, the next two lower lines being partly blended with lines of unrelated species. The average spectrum obtained toward IRC+10216 is plotted in Figure 2a. It has an r.m.s. noise per 1 MHz channel of 0.90 mK in the $T_A^*$ scale.

The carbon-chain molecules observed in IRC+10216 are concentrated in a thin shell of radius $\simeq 15''$ (see Fig.2b). Observed with the 30-m telescope, the profiles of their $\lambda 3$ mm lines have the same cusped shape and the same width (29.5 kms$^{-1}$). They are all centred at $V_{LSR} = -26.5$ kms$^{-1}$. The spectrum of Fig. 2a shows two cusped lines, near 89.75 GHz and three weaker features near 89.80 GHz. The former can be readily assigned to the $N = 4 \rightarrow 3$, $F = 4.5 \rightarrow 3.5$ and $F = 3.5 \rightarrow 2.5$ fine structure components of C$_3$H in its first excited bending state, $\nu_4 = 1^2 \Sigma^u$ (Yamamoto et al. 1990) since they have
the right frequencies and the right intensity ratio (1.33±1, whereas the expected LTE ratio is 1.30).

We have fitted 3 cusped lines (assuming standard line shapes and 29.5 kms\(^{-1}\) widths) to the 3 weak spectral features of Fig. 2a. The derived intensities and rest frequencies are given in Table 1; the fitted spectrum is compared in Fig. 2c to the observed spectrum. The first two lines agree in frequency with the \((N, J) = (32, 32.5) \rightarrow (31, 31.5)\) and \((32, 31.5) \rightarrow (31, 30.5)\) transitions of C\(_5\)N and very probably arise from this radical. Indeed, although the \(\lambda 3 \text{ mm}\) spectrum of IRC+10216 is more crowded than the 1 cm spectrum of TMC1, there are not many lines that we cannot assign to the rotational transition of a known circumstellar molecule: in our 30-m telescope spectral survey of IRC+10216, the density of unidentified lines stronger than \(\geq 3 \text{ nK}\) is only of 1 per 80 MHz (see Guélin et al. 1997). The probability that two unrelated lines lie within 1 MHz from the C\(_5\)N lines is thus lower than 1 per one thousand. Whereas the first two features can be tentatively assigned to C\(_5\)N, the third, which is weaker and lies 7 MHz higher in frequency, remains unidentified.

### 3. The abundance of C\(_5\)N

In the direction of the TMC 1 cyanopolyne peak, the rotational populations of HC\(_5\)N and HC\(_5\)N can be described by Boltzmann distributions with rotation temperatures of \(T = 10 K\) (see e.g. Takano et al. 1997). We adopt therefore \(T_{rot} = 8 K\). The H\(_2\) column density in this direction is \(N(H_2) = 10^{22} \text{ cm}^{-2}\) (Cernicharo & Guélin 1987). That of C\(_5\)N can be calculated from the standard expression for the optically thin lines of thermalized linear molecules:

\[
N(C_5N) = 0.7 \times 10^{16} \frac{T_{rot} T'_0}{T_{rot} - T'_{bg}} (\mu \nu)^{-2} e^{-E_u/kT_{rot}} \Sigma \int T_{MB} dv,
\]

where

\[
T' = (h^2/k)(e^{\frac{E_u}{kT_{rot}}} - 1)^{-1},
\]

and where \(\Sigma \int T_{MB} dv = 0.014 \text{ Kkms}\(^{-1}\) is the sum of the integrated intensities of the two doublet components. In equation [1], \(N\) is in \(\text{cm}^{-2}\), the dipole moment \(\mu_0 = 3.385 \text{ D}\) (Botschwina 1996) in debye, and the line frequency \(\nu = 25.25 \text{ GHz}\) in gigahertz. We find:

\[
N(C_5N) = 3.1 \times 10^{11} \text{ cm}^{-2},
\]

\[
x(C_5N) = N(C_5N)/N(H_2) \approx 3 \times 10^{-11}.
\]

The abundance of C\(_5\)N can be compared to those of the related species HC\(_5\)N and C\(_5\)N in the light of the chemical model predictions. We calculate first the abundance of the \(^{13}\text{C}\) isotopomers of HC\(_5\)N, whose 1 cm lines are optically thin, by setting in equation [1] \(\mu_0 = 4.33 \text{ D}, \nu = 23.7 \text{ GHz}\) and \(\int T_{MB} dv = 0.065 \text{ Kkms}\(^{-1}\), which is the average of the integrated intensities of the J=9--8 lines of HC\(_{13}\)C\(_5\)N and HC\(_4\)\(^{13}\)CN, observed with the Effelsberg telescope (Takano et al. 1998). We find \(N(\text{HC}_{13}\text{CN}) = 1.0 \times 10^{12} \text{ cm}^{-2}\), from which we derive \(N(\text{HC}_5\text{N}) = 7 \times 10^{13} \text{ cm}^{-2}\), adopting the

‘s’standard’ elemental abundance ratio \(^{12}\text{C}/^{13}\text{C} \approx 70\) in the local interstellar medium (see Wilson & Rood 1994). The value of \(7 \times 10^{13} \text{ cm}^{-2}\) is close to the value of Suzuki et al. (1992), as well as to the value we estimate with an LVG code from the main isotopomer \(J = 9 \rightarrow 8\) line intensity (5 \times 10^{13}\text{cm}^{-2}). We arrive at:

\[
N(\text{HC}_{13}\text{CN}/C_5N) = 3
\]

\[
N(\text{HC}_5\text{N}/C_5N) \approx 200.
\]

For C\(_5\)N, we find, using in eq. [1] \(T_{rot} = 8K, \mu_0 = 2.84 \text{ D}\) (Pauzat et al. 1991), and the line parameters of Table 1: \(N(C_5N) = 8.2 \times 10^{12} \text{ cm}^{-2}\). Finally, we take for HC\(_5\)N the column densities derived by Takano et al. (1998),

\[
N(\text{HC}_{13}\text{CCN}) = 2.1 \times 10^{12} \text{ cm}^{-2}\text{ and N(}^1\text{H}12\text{C}_3\text{N}) = 1.6 \times 10^{14} \text{ cm}^{-2}.
\]

This yields:

\[
N(\text{HC}_{13}\text{CCN}/C_5N) = 0.26
\]

\[
N(\text{H}12\text{C}_3\text{N}/C_5N) = 19.
\]

The C\(_5\)N/HC\(_5\)N abundance ratio is an order of magnitude smaller than the C\(_5\)N/HC\(_5\)N ratio. According to neutral-neutral gas phase chemical models, the cyanopolyne mole-cules are mainly formed by reactions of N and CN with polyacetylenes or polyacetylic ions (Herbst & Leung 1990). The cyanopolyne radicals are formed in TMC1 by the reaction of atomic C with cyanopolyynes (Herbst et al. 1994), and in IRC+10216 by photodissociation (Cherchneff & Glassgold 1993). They are destroyed by reactions with N atoms, O atoms, polyacetylenes and photodissociation.

According to model predictions, HC\(_5\)N/HC\(_5\)N varies in TMC1 by several orders of magnitude between the “early times” and steady state. The C\(_5\)N/HC\(_5\)N and C\(_5\)N/HC\(_5\)N ratios, on the other hand, remain constant within a factor of 2. They are comprised between 0.1 and 0.2 (see Table 3 of Herbst et al. 1994). Whereas the observed C\(_5\)N/HC\(_5\)N ratio (0.2, Cernicharo et al. 1987) agrees with the predicted one, C\(_5\)N/HC\(_5\)N is more than one order of magnitude too low. The most recent models, which take into account the destruction of acetylene and polyacetylenes by C atoms and the reaction of CN with O, form enough C\(_5\)N but too little C\(_5\)N. Here also, C\(_5\)N is underabundant with respect to model predictions (Cherchneff & Glassgold 1993), and C\(_5\)N/HC\(_5\)N one order of magnitude smaller than C\(_5\)N/HC\(_5\)N.

In the case of IRC+10216, we adopt for C\(_5\)N the same rotation temperature as for HC\(_5\)N (\(T_{rot} = 29 K\), Kawaguchi et al. 1996). We then derive from the integrated intensities of Table 1 a line-of-sight column density in the direction of the central star (twice the radial column density across the shell) \(N(C_5N) = 6.1 \times 10^{12} \text{ cm}^{-2}\). This is ~ 50 times less than the column densities of C\(_5\)N and HC\(_5\)N. Here also, C\(_5\)N is underabundant with respect to model predictions (Cherchneff & Glassgold 1993), and C\(_5\)N/HC\(_5\)N one order of magnitude smaller than C\(_5\)N/HC\(_5\)N.

The unexpectedly low C\(_5\)N/HC\(_5\)N abundance ratio in found both sources shows that the formation of long carbon-chain molecules is not fully understood, and that is difficult to predict the abundances of unobserved species. The very long carbon chains could be more abundant than
Table 1: Observed line parameters

| Rest. frequ. (MHz) | Obs.-Calc. (MHz) | Transition $N, J \rightarrow N', J'$ | Species | $\int T_{MB}dv$ (mK km s$^{-1}$) |
|-------------------|-----------------|-------------------------------------|--------|---------------------|
| TMC1              |                 |                                     |        |                     |
| 19799.956         | .005            | 2.1,5.1-1,0.5                      | C$_3$N | 23 (8)              |
| 19800.121         | .000            | 2.1,5.1-1,0.5                      | C$_3$N | 58 (8)              |
| 19780.801         | .001            | 2.2,5.1-1,1.5                      | C$_3$N | 60 (8)              |
| 19780.826         | .000            | 2.2,5.1-1,1.5                      | C$_3$N | 34.5 (8)            |
| 19781.096         | .002            | 2.2,5.5-1,1.5                      | C$_3$N | 86 (8)              |
| 23963.897         | 9.000           | 9-8                                | HC$_3$N| 2240 (100)          |
| 25249.938 (4)     | .018 (40)       | 9.9-5-8.8,5                        | C$_5$N | 7.3 (9)             |
| 25260.649 (4)     | .017 (40)       | 9.8-5-7.5                          | C$_5$N | 6.4 (9)             |
| IRC+10216         |                 |                                     |        |                     |
| 89730.54(10)      | -.07 (10)$^b$   | 4.45 - 3.35                         | C$_3$H($v^2\Sigma^u_a$) | 880(20) |
| 89759.17(12)      | -.18 (12)$^b$   | 4.35 - 3.25                         | C$_3$H($v^2\Sigma^u_a$) | 660(20) |
| 89785.6 (4)       | .4 (13)         | 32,32.5-31,31.5                     | C$_3$N | 95 (15)             |
| 89797.0 (3)       | 1.1(13)         | 32,31.5-31,30.5                     | C$_5$N | 105(20)             |
| 89804.0 (10)      | -               | -                                   | U     | 75 (20)             |

Notes to the Table:

$^b$: weighted average of the two blended hyperfine components. The calculated frequencies are taken from Kasai et al. (1997) for C$_3$N, Yamamoto et al. (1990) for C$_3$H, and Guélin et al. (1982) for C$_5$N (see also Gottlieb et al. 1983). The observed rest frequencies of C$_3$N and C$_5$N were derived assuming $V_{LSR} = 5.65$ kms$^{-1}$ in TMC 1 and -26.5 kms$^{-1}$ in IRC+10216; that of HC$_3$N was derived assuming $V_{LSR} = 5.75$ kms$^{-1}$ (see Guélin et al. 1982). The number in parenthesis represent the r.m.s. uncertainty on the last digit. The uncertainties on the line integrated intensities, given in the Table, do not include the calibration uncertainty which is 10 – 20%. Note that the intensity ratio between the rotational transitions in the C$_3$H ground state and in the $^2\Sigma^u$ excited bending state, hence presumably the population ratio between these two states, is $\sim 10$.

Expected. We note, however, that the species of medium size, such as the chains consisting of 4–7 C, N, or O atoms and the rings with less than 10 heavy atoms, which would give rise in TMC 1 to a rich centimetric spectrum, are probably not very abundant in that source: we have covered so far a 100 MHz-wide band in TMC 1 with a very good sensitivity and did not detect any unidentified line down to a level of 10 mK. The acetylene chains and cumulene carbenes appear in this respect exceptional.

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