

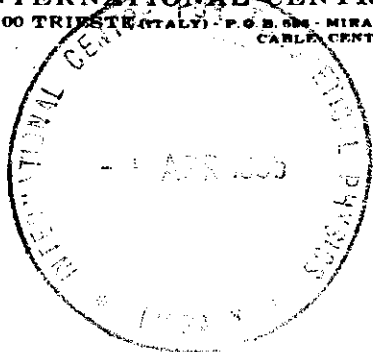


INTERNATIONAL ATOMIC ENERGY AGENCY
UNITED NATIONS EDUCATIONAL, SCIENTIFIC AND CULTURAL ORGANIZATION



INTERNATIONAL CENTRE FOR THEORETICAL PHYSICS

34100 TRIESTE (ITALY) - P.O. BOX - MIRAMARE - STRADA COSTIERA 11 - TELEPHONE: 0431/574466
CABLE CENTRATON - TELEX 480382-I



SMR/115 - 57

WINTER COLLEGE ON LASERS, ATOMIC AND MOLECULAR PHYSICS

(21 January - 22 March 1985)

INTERSTELLAR MOLECULES

P. ENCRENAZ
Meudon Observatoire
92190 Meudon
France

These are preliminary lecture notes, intended only for distribution to participants.
Missing or extra copies are available from Room 229.

Introduction:

Advances in spectroscopy using large optical telescopes has led to the discovery of a fair amount of

— diffuse band in the optical region (1940-20) which have not been interpreted until now, and may have been identified in 1984-5 by Legen and coworkers

— narrow absorption lines (1937-1943) in the spectra of stars by Smith, Adams and Swings. These lines have been attributed to the CN, CH and CH⁺ radicals (Herbig, Swings), but were not used until 1968 to measure the 3K background radiation (discovered by Penzias^{Wilson} in 1964-5 and measured directly with the horn antenna)

Shklovsky and Townes suggested to look for OH in the late fifties, but OH was not detected until 1963 in absorption in front of Cas A. In 1968 H₂O and NH₃ were detected by Townes and his collaborators, while H₂CO, a fragile molecule in the interstellar environment, could not do better than push for a thorough search for interstellar molecules in the millimeter region of the electromagnetic spectrum. CO, CN, CH₃CN were promptly observed using a sensitive heterodyne millimeter receiver, and the field has now exploded since more than 60 molecules and 1000 transitions have been observed as of today.

A new field has been opened, which needs a strong coupling between spectroscopy, collisions in atomic and molecular physics, radioastronomy, chemistry.

$$T_{ex} = 2.8 \pm 0.1 \text{ K}$$

218

- Galactic and Extra-Galactic Radio Astronomy

be obtained under any other reasonable assumption. While this does not prove the existence of the 3°K background, it corroborates the firmer evidence which other molecules are able to provide.

9.3.3 Optical Molecules and the 3°K Cosmic Radiation

An interesting feature of most of the light diatomic molecules observed in the interstellar medium at optical wavelengths is that their ground-state rotational energy level schemes are characterized by $E/k \sim h^2/2Ik \sim 3^\circ\text{K}$ (I is the moment of inertia of the molecule). Hence these molecules are

extremely sensitive indicators of the density of the 3°K field. Figure 9.2 shows observed spectra and relevant energy levels of CH, CH⁺, and CN used for sampling the 3°K field. The relative intensities of the lines shown depend on the temperature of the radiation field, assuming that the rotational level populations are in equilibrium with this field. Those lines which are not observed [R(2) in CN, R(1) in CH⁺, and R₁(1) in CH] indicate that only the lowest levels are populated under interstellar conditions. The energy levels show that CN samples the radiation field at 2.64 and 1.32 mm by means of the intensity ratios R(0)/R(1) and R(2)/R(1); similarly CH⁺ samples the field at 0.359 mm

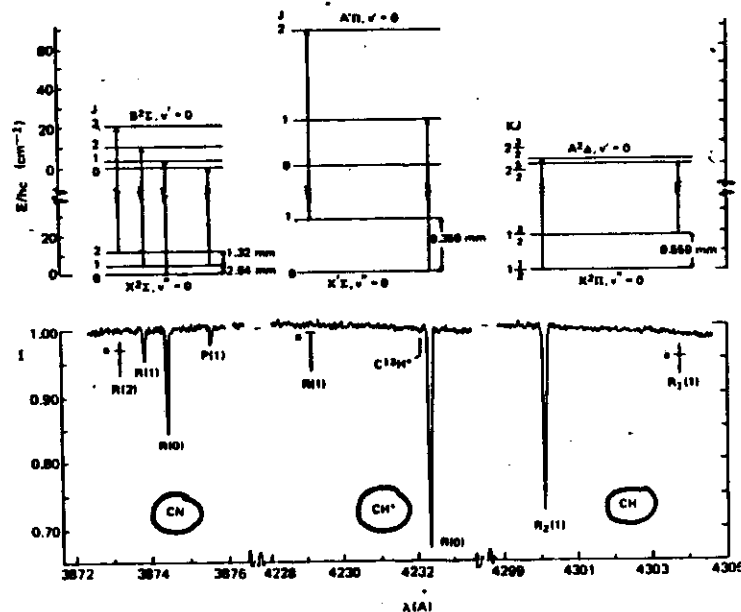


Figure 9.2 Observed spectra in the UV of the interstellar molecules CN, CH⁺, and CH. The observed transitions are indicated on the energy level diagrams, as are the mm-wavelength transitions whose excitation temperatures are inferred to be that of the cosmic background radiation. (Figure reproduced by kind permission of P. Thaddeus.)

Intensité relative des transitions élect. → températures d'excitation des niveaux de rotation

②

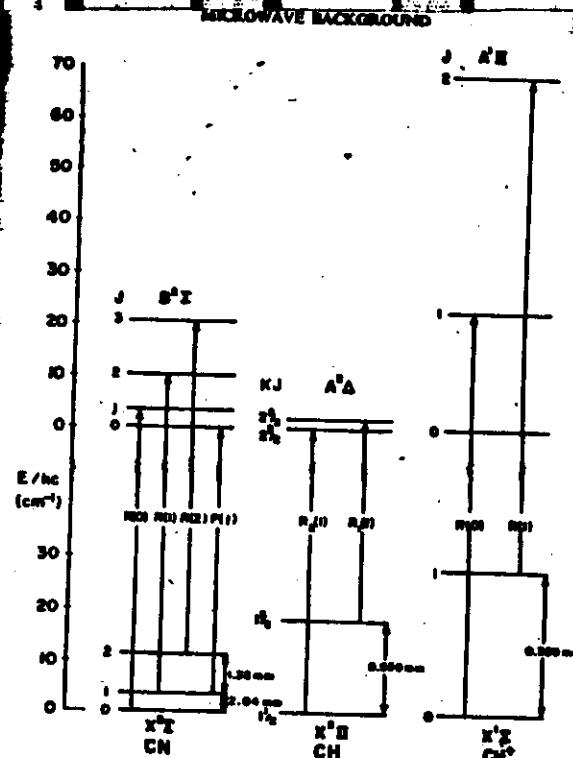


Figure 3. Term diagram of the strongest interstellar bands of CN, CH, and CH⁺. All levels are in the lowest vibrational state.

extract from the molecular spectra extremely valuable upper limits to the radiation intensity at wavelengths even shorter than 2.64 mm. It is therefore appropriate to consider in some detail the current state of these observations, and also the theoretical analyses of local processes which show that the molecules are indeed reliable thermometers for the background radiation.

③

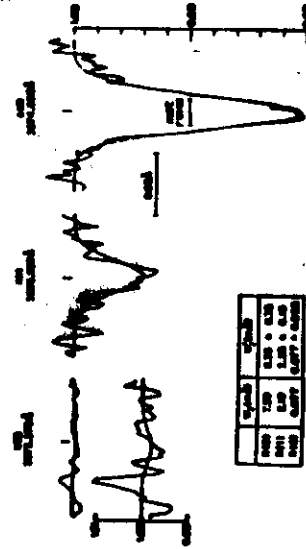


Figure 1. The microwave CH line in f O₂ observed with a high-resolution Fourier transform spectrometer (Nagel, Thiele & Carlson 1972). The region of R23 was scanned for many nights, that of R20 and R21 for much shorter times.

Since we will have presented as though the electronic ground state were ²Σ, and the level structure, as shown in Figure 3, that of a simple rigid rotor. Since only the two lowest rotational levels are observed to be populated in space, the thermal level populations n_0 , n_1 in statistical equilibrium are determined from the rate equation

$$\frac{dn_1}{dt} = -n_1(C_{10} + A_1) + n_0 C_{01} = 0$$

where A_1 is the spontaneous relative decay rate of the $J=1$ level,

$$A_1 = \frac{64\pi^2 \nu^3}{3c^3} = 1.10 \times 10^{-4} \text{ sec}^{-1}$$

$\nu = 1.40 \times 10^{14} \text{ cm}^{-1}$ is the CH dipole moment (Thomson & Dalby 1969) and $n_0 = 1.13 \times 10^{10} \text{ cm}^{-3}$, C_{01} is the direct rate of radiative or collisional de-excitation from the $J=1$ to $J=0$ level, and C_{10} is the rate of excitation from the $J=0$ level to all others, since excitation to a level above $J=1$ is in effect immediately followed by relative cascade to that level. Under neutral particle impact the cross sections for many-quantum transitions will be comparable to those for the $\Delta J=1$ transitions, and C_{01} and C_{10} as defined will be suitably doubled between.

$$\frac{C_{01}}{C_{10}} = \frac{n_1}{n_0} \exp(-h\nu/kT) \approx \frac{n_1}{n_0}$$

where $n_1/n_0 \approx 1$, and T is the kinetic temperature. But since the cross sections for neutral particle excitation can only be estimated, while those for excitation

by H₂ or He are rather irregularly known, and only at low energies the selection rule $\Delta J=1$, it is preferable in the treatment of space to assume that Equation 3 is satisfied. Then the optical observation of neutral CH requires

$$n_1/n_0 = (n_1/n_0) \exp(-h\nu/kT), \text{ with } T_0 = 2.5\% \quad 4$$

which may be solved for the excitation rate, yielding

$$C_{01} = (n_1/n_0) A_1 \exp(-h\nu/kT) [1 - \exp(-h\nu/kT)]^{-1} \quad 5$$

$$= 0.0 \times 10^{-4} \text{ sec}^{-1}$$

It is the rate required to account for the observed CH emission, and it is the rate obtained from 2.5% excitation.

The excitation equation we now wish to answer is whether excitation at the rate C_{01} can be provided by any process in the CH cloud besides direct photo-excitation. We shall consider additional input of the CH molecules with the $J=1$ ground state, either (a) with the most abundant neutral particles, presumably H₂ or He, or (b) with the excited H I atoms; or (c) with the excited H₂ molecules, or (d) with the excited He atoms. Since the CH molecules, or He atoms, are not expected, in the edge of an ionization front and are subject to a $\sim 10\%$ protonic, and doubly (or) molecular of the rotational level is optical photo-ionization (protonic ionization). Ionization via the vibration of CH, either by collision or infrared photons, is such a slow process that it is not to be considered.

Neutral particle impact.—Rotational excitation by H₂ or He, impact is a complex collisional process dependent on short-range intermolecular forces, but by the time the excitation cross section is known, it is not possible to estimate it. It is reasonable to adopt an upper limit $\sigma \sim 2 \times 10^{-20} \text{ cm}^2$. Taking $n_0 \sim 10^{10} \text{ cm}^{-3}$ as an upper limit to the density in the f O₂ molecular cloud, and $n_1 \sim 1.2 \times 10^{10} \text{ cm}^{-3}$ (i.e., 10% H₂ atoms) as an upper limit to the kinetic velocity, yields

$$C_{01} \text{ as upper } 5.2 \times 10^{-4} \text{ sec}^{-1} \quad 6$$

which falls short of Equation 5, the required rate, by a factor of about 25. The density of molecular hydrogen, for which the excitation cross section is comparable to that of H₂, is unknown in the f O₂ cloud, but it is unlikely to exceed the density of the H atoms. Almost certainly, therefore, neutral particle impact is not the cause of the CH excitation. There is other evidence pointing to this conclusion: it is the density required for sub-application of the CO excitation temperature (Schuler & Stahler 1971) and probably the optical excitation would be much larger than observed.

Electron impact.—Because of the long range of the Coulomb force, electron impact at H I energies is highly effective in exciting rotational levels of polar molecules, its rate coefficient being ~ 10 times that of neutral particles. Excitation by the thermal electrons in an H I region is for this reason the least process which seems closest to accounting for the 2.5% CH excitation, and it accordingly de-

4

5

These molecules, whose list is given in table I, are made of the most abundant elements (H, C, N, O, S, Si, Fe), and their isotopic substitutes are also in detail (H/D, ¹²C/¹³C, ¹⁴N/¹⁵N, ¹⁶O/¹⁷O/¹⁸O, ²⁸Si/²⁹Si/³⁰Si, ³²S/³³S/³⁴S

Before we try to understand the impact of these discoveries on astronomy, I will go back to basic and simple principles for the excitation of these molecules, and I will spend some time on the case of neutral atomic hydrogen (H I).

Table of the most abundant elements

H	3.18	10 ¹⁰
C	4.18	10 ⁹
N	3.7	10 ⁶
O	2.15	10 ³
S	5	10 ⁵
Si		10 ⁶
Cl	5.7	10 ³
Fe	6.3	10 ⁵

Isotopic Ratios (see neighborhood)

H/D	5 x 10 ⁵
¹² C / ¹³ C / ¹⁴ C	89 / radioactive
¹⁴ N / ¹⁵ N	280
¹⁶ O / ¹⁷ O / ¹⁸ O	3000 / 500
³² S / ³³ S / ³⁴ S	100 / 20
²⁸ Si / ²⁹ Si / ³⁰ Si	20 / 25
³⁵ Cl / ³⁷ Cl	4 / 1

6

des des riles dans l'atmosphère qui les microscopiques ont observés pour H₂, CO et d'autres molécules dans le milieu interstellaire et les enveloppes d'étoiles froides. Ce sont cependant les transitions de rotation, qui sont les plus nombreuses radio centimétriques, millimétriques et submillimétriques qui ont permis la plupart des découvertes de molécules interstellaires. Les premiers abaissements d'énergie de rotation des molécules correspondants à des énergies peu élevées et les collisions entre les molécules et les particules de milieux interstellaires sont facilement leur réaction : des réactions rapides se déroulent dans les photochimie et les réactions correspondantes. C'est ainsi qu'a été découverte la très grande majorité des molécules interstellaires et circumstellaires (celles qui se trouvent dans les enveloppes des étoiles). L'analyse donne la liste des molécules interstellaires et circumstellaires.

La plupart des molécules détectées sont linéaires : les atomes qui les constituent se succèdent les uns aux autres le long d'une ligne droite ou brisée, non fermée sur elle-même (ce qui définirait une molécule cyclique). L'exemple le plus simple des acides aminés, à cet égard, est le glycine.

Les molécules linéaires sont les plus nombreuses dans les nuages interstellaires. Les autres qui les constituent se succèdent les uns aux autres le long d'une ligne droite ou brisée, non fermée sur elle-même (ce qui définirait une molécule cyclique). L'exemple le plus simple des acides aminés, à cet égard, est le glycine.

Ce milieu se caractérise également par la présence de poussières interstellaires, grains minuscules de silicate et de graphite de tailles inférieures au micromètre, mais aussi moléculaires en gaz. Elles sont nombreuses et leur masse totale atteint 1 à 2 % de celle du gaz. D'une part, dans le verre, elles constituent un site privilégié pour la formation d'une partie des molécules ; d'autre part, elles absorbent et diffusent la lumière des étoiles, si bien que les régions centrales d'un nuage ne sont éclairées que par la lumière diffusée par les grains et sont donc privées de la lumière des étoiles. Elles jouent également un rôle important dans la destruction des molécules. Dans ce cas, et par conséquent, les régions les plus denses où se produisent les réactions sont dépourvues de molécules, on a affaire à ce qu'on appelle un milieu interstellaire.

et O. On trouve aussi des molécules circumstellaires et des étoiles, en fait, on trouve des molécules dans les nuages interstellaires. Les premières ont été découvertes par la spectroscopie infrarouge, c'est-à-dire par l'analyse des raies d'émission des étoiles. Les autres ont été découvertes par la spectroscopie radio, c'est-à-dire par l'analyse des raies d'émission des étoiles.

Pour former une molécule, les atomes doivent d'abord se rencontrer. Les premières ont été découvertes par la spectroscopie infrarouge, c'est-à-dire par l'analyse des raies d'émission des étoiles. Les autres ont été découvertes par la spectroscopie radio, c'est-à-dire par l'analyse des raies d'émission des étoiles.

Dans un tel milieu, la chimie est très simple. Les premières ont été découvertes par la spectroscopie infrarouge, c'est-à-dire par l'analyse des raies d'émission des étoiles. Les autres ont été découvertes par la spectroscopie radio, c'est-à-dire par l'analyse des raies d'émission des étoiles.

LES MOLECULES INTERSTELLAIRES OU CIRCUMSTELLAIRES CONNUES A CE JOUR

2 atomes	3 atomes	4 atomes	5 atomes	6 atomes	7 atomes	8 atomes	9 atomes	10 atomes	11 atomes	12 atomes	13 atomes	14 atomes	15 atomes	16 atomes	17 atomes	18 atomes	19 atomes	20 atomes	21 atomes	22 atomes	23 atomes	24 atomes	25 atomes	26 atomes	27 atomes	28 atomes	29 atomes	30 atomes	31 atomes	32 atomes	33 atomes	34 atomes	35 atomes	36 atomes	37 atomes	38 atomes	39 atomes	40 atomes	41 atomes	42 atomes	43 atomes	44 atomes	45 atomes	46 atomes	47 atomes	48 atomes	49 atomes	50 atomes	51 atomes	52 atomes	53 atomes	54 atomes	55 atomes	56 atomes	57 atomes	58 atomes	59 atomes	60 atomes	61 atomes	62 atomes	63 atomes	64 atomes	65 atomes	66 atomes	67 atomes	68 atomes	69 atomes	70 atomes	71 atomes	72 atomes	73 atomes	74 atomes	75 atomes	76 atomes	77 atomes	78 atomes	79 atomes	80 atomes	81 atomes	82 atomes	83 atomes	84 atomes	85 atomes	86 atomes	87 atomes	88 atomes	89 atomes	90 atomes	91 atomes	92 atomes	93 atomes	94 atomes	95 atomes	96 atomes	97 atomes	98 atomes	99 atomes	100 atomes
----------	----------	----------	----------	----------	----------	----------	----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	-----------	------------

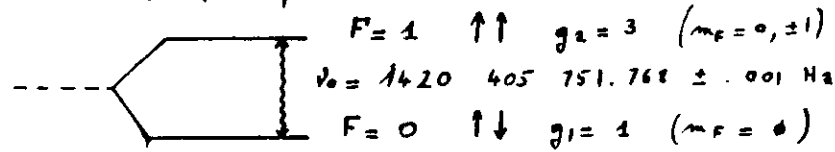
Remarque : les molécules de 2 atomes sont les plus nombreuses, suivies de celles de 3 atomes, puis de celles de 4 atomes, etc. Les molécules de 10 atomes ou plus sont très rares.

(1) P. Thaddeus, I.E. Charnick, A.A. Lutz, Astrophysical Journal Letters, 1961.

7

A] Atomic Neutral Hydrogen: HI

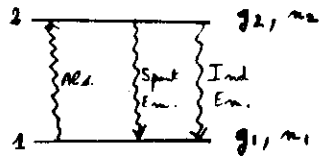
① The 21 cm line was discovered in 1951. This transition connects two hyperfine levels of atomic hydrogen with either parallel or antiparallel spins of the proton and electron.



The spontaneous transition rate $A_{21} = 2.84 \times 10^{-15} \text{ s}^{-1}$ is exceedingly small (magnetic dipole transition). The lifetime of the excited atom is therefore of the order of 10^7 years.

A] ② Line formation:

Let $I(\nu)$ be the specific intensity of the radiation field at the frequency ν : $I(\nu) = \frac{c}{4\pi} u(\nu)$ ($u(\nu)$ in $\text{erg cm}^{-3} \text{ Hz}^{-1}$ = spectral density energy) = $\text{erg cm}^{-2} \text{ s}^{-1} \text{ Hz}^{-1}$



By definition, the Einstein coefficients are given by:

$$g_1 B_{12} = g_2 B_{21}$$

$$B_{21} = \frac{c^2}{2h\nu^3} A_{21}$$

with $N_{1 \rightarrow 2} = n_1 B_{12} I(\nu)$

$$N_{2 \rightarrow 1} = n_2 (A_{21} + B_{21} I(\nu))$$

(number of transition $1 \rightarrow 2$ and $2 \rightarrow 1$ per unit time and unit volume).

If $I(\nu) = 0$, the spontaneous lifetime gives $\tau_2 \approx \frac{1}{A_{21}}$ (coupling with

② the fluctuation of the vacuum).

The variation of $I(\nu)$ along an elementary path ds can therefore be written $\frac{dI(\nu)}{ds} = \frac{1}{8\nu} \times \frac{h\nu}{4\pi} \times (N_{2 \rightarrow 1} - N_{1 \rightarrow 2})$ as all photons are redistributed in a frequency interval $\delta\nu$.

The equation is usually written $\frac{dI(\nu)}{ds} = j(\nu) - \kappa(\nu) I(\nu)$

$$j(\nu) \text{ in } \text{erg cm}^{-1} \text{ s}^{-1} \text{ Hz}^{-1}$$

$$\kappa(\nu) \text{ in } \text{cm}^{-1}$$

By definition the optical depth is $d\tau(\nu) = -\kappa(\nu) ds$

$$\kappa(\nu) = \frac{1}{8\nu} \frac{h\nu}{4\pi} n_1 \frac{g_2}{g_1} \frac{c^2}{h\nu^3} A_{21} \left(1 - \frac{n_2}{n_1} \frac{g_1}{g_2}\right)$$

Or, if we introduce the excitation temperature T_{ex} such that

$$\frac{n_2}{n_1} = \frac{g_2}{g_1} \exp\left(-\frac{h\nu}{kT_{ex}}\right)$$

$$\kappa(\nu) = \frac{1}{8\nu} \frac{h\nu}{4\pi} n_1 \frac{g_2}{g_1} \frac{c^2}{2h\nu^3} \left(1 - \exp\left(-\frac{h\nu}{kT_{ex}}\right)\right)$$

or, if $h\nu \ll kT_{ex}$

$$\kappa(\nu) = \frac{1}{8\nu} \frac{h\nu}{4\pi} n_1 B_{12} \frac{h\nu}{kT_{ex}}$$

③ Population of the levels 1 and 2:

The 2 level system is supposed to interact only with the radiation field (characterised by $I(\nu)$) and with particles (inelastic collisions). These particles (H, H₂, ions...) have a Maxwellian Distribution of kinetic temperature T_k .

$$f(u) du = \left(\frac{m}{2\pi kT_k}\right)^{3/2} e^{-\frac{mu^2}{2kT_k}} 4\pi u^2 du$$

m reduced mass

The population of levels 1 and 2 changes with T_k , but we will assume that the equilibrium is reached before $I(\nu)$ and T_k change.

significantly:

$$\frac{dn_1}{dt} = \frac{dn_2}{dt} = 0$$

(10)

$$n_1 B_{12} I(\nu) + n_1 C_{12} = n_2 B_{21} I(\nu) + n_2 A_{21} + n_2 C_{21}$$

C_{12} (and C_{21}) are the probabilities of collisional excitation, and de-excitation)

It can be shown that the Maxwellian hypothesis imply $n_1 C_{12} = n_2 C_{21}$ and that $T_{12} = T_K$. We have thermodynamic equilibrium.

[R] if $\frac{dI}{dt} = 0$ (system isolated from all external radiation), then

$$I(\nu) = \frac{j(\nu)}{\kappa(\nu)} = \frac{2h\nu^3}{c^2} \frac{1}{\left[\exp\left(\frac{h\nu}{kT}\right) - 1\right]} \quad (\text{black body spectrum})$$

$$\frac{n_2}{n_1} = \frac{B_{12} I(\nu) + C_{12}}{B_{21} I(\nu) + A_{21} + C_{21}}$$

Two interesting limiting cases: - $C_{21} \gg A_{21} \left[1 + I(\nu) \frac{c^2}{2h\nu^3} \right]$
and and and $T_{12} = T_K$

$$- C_{21} \ll A_{21} \left[1 + I(\nu) \frac{c^2}{2h\nu^3} \right]$$

$$T_{12} = T_K$$

④ 21 cm transition: $\frac{h\nu}{k} = .07 \text{ K}$

$$T_K \sim 100 \text{ K}$$

$$T_R = 2.8 \text{ K (cosmological background)}$$

$$\frac{c^2}{2h\nu^3} I(\nu) = 43.6$$

$$\nu = 9.8 \times 10^3 \sqrt{T_K} \text{ cm}^{-1}$$

$$\sigma_{\text{HI}} \sim 10^{-18} \text{ cm}^2$$

Collisions are dominant if $n > 10^{-3} \text{ cm}^{-3}$. This is ALWAYS the case

$$\begin{array}{|c|} \hline T_{12} = T_K \\ \hline \end{array}$$

$$\rightsquigarrow I(\nu)$$

A photon of frequency ν will interact only with the class of HI atoms having the velocity v such that $\frac{v}{c} = \frac{\nu - \nu_0}{\nu_0}$ (within the natural width of the line).

The transfer equation can be solved and give

$$I(\nu) = I_0(\nu) \exp[-\tau(\nu)] + B(\nu) [T_{12}] [1 - \exp(-\tau(\nu))]$$

$$\tau(\nu) = \int_L \kappa(\nu) ds$$

$$B(\nu) = \frac{2h\nu^3}{c^2} \left[\exp\left(\frac{h\nu}{kT_{12}}\right) - 1 \right]^{-1} = \frac{j(\nu)}{\kappa(\nu)}$$

By definition the brightness temperature T_B is the temperature of a black body having $I(\nu)$ at frequency ν . Using the Rayleigh - Jeans approximation ($B(\nu) = \frac{2kT\nu^2}{c^2}$), the equation becomes

$$T_B(\nu) = T_{B0}(\nu) e^{-\tau(\nu)} + T_{12} (1 - e^{-\tau(\nu)})$$

$\tau(\nu) = 5.48 \times 10^{-14} \frac{N_{\text{HI}} T_K}{T}$ ($P(\nu)$ is the probability of the HI atom to have a projected velocity between v and $v+dv$).

Observations: - on a cloud at two frequencies

$$T_B^{\text{obs}}(\nu) = T_{12} \tau(\nu) + T_{B0}(\nu)$$

$$T_B^{\text{obs}}(\nu') = T_{B0}(\nu') = T_{B0}(\nu)$$

$$\Delta T_B \sim 5.48 \times 10^{-14} N_{\text{HI}} P(\nu)$$

Si on somme sur la raie

$$\int_{\text{raie}} \Delta T_B(\nu) d\nu = 5.48 \times 10^{-14} N_{\text{HI}}$$

$$N_{\text{HI}} = 1.8 \times 10^{18} \int_{\text{raie}} \Delta T_B d\nu \text{ cm}^{-2}$$

- on and off a cloud at the same frequency ⁽¹²⁾

$$T_B^{\text{on}} = T_{B_0} e^{-\tau(\nu)} + T_{\text{in}} (1 - e^{-\tau(\nu)})$$

$$T_B^{\text{off}} = T_{\text{in}} (1 - e^{-\tau(\nu)})$$

$$\Delta T_B(\nu) = T_{B_0} [1 - e^{-\tau(\nu)}] \Rightarrow \text{measure of } \tau(\nu)$$

The best use is the use of pulsars for measuring $\tau(\nu)$ and the distance of these objects.

B] The case of simple molecules: ⁽¹³⁾

If you lower the temperature of a cloud of neutral hydrogen, its density will increase and you will form molecules. The most abundant of these molecules, H_2 , cannot be detected by its microwave spectrum. But CO, the second most abundant, has such a simple spectrum that it is quite straightforward to understand it completely:

a) linear molecule: $E(J)/k = B_0 J(J+1) - D_J [J(J+1)]^2$

The centrifugal distortion is 10^{-4} to 10^{-6} smaller than the rotational term.

$$\nu(J, J-1) \approx 2B_0 J$$

	B_0 (MHz)	D MHz	μ Debye	J	$A_{J, J-1}^{(s-1)}$
^{12}CO	57636	183	0.11	1	6×10^{-8}
^{13}CS	24496	40	1.95	3	6×10^{-5}
HCO^+	44594	.09	4.02	1	5×10^{-5}

$$A_{J, J-1} = \frac{64 \pi^4 \mu^2}{3 h c^3} \nu^3 \frac{J}{2J+1}$$

$$A_{J, J-1} = 9.3 \cdot 10^{-20} B_0^3 \mu^2 \frac{J^4}{2J+1} (\text{s}^{-1}) \left(\frac{B_0}{\text{MHz}} \right) \left(\frac{\mu}{\text{Debye}} \right)^2 (10^{-18} \text{ erg})$$

Certain molecules are smart entities you can see millions of times

If we follow the same reasoning then in A], we obtain

$$g_{J-1} C_{J-1, J} = g_J C_{J, J-1} \exp\left(-\frac{h\nu}{k T_K}\right)$$

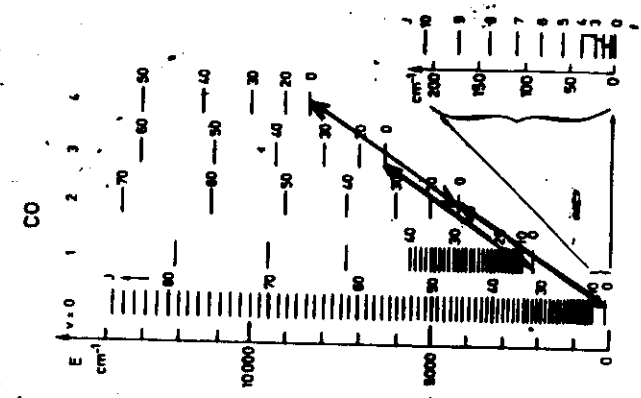
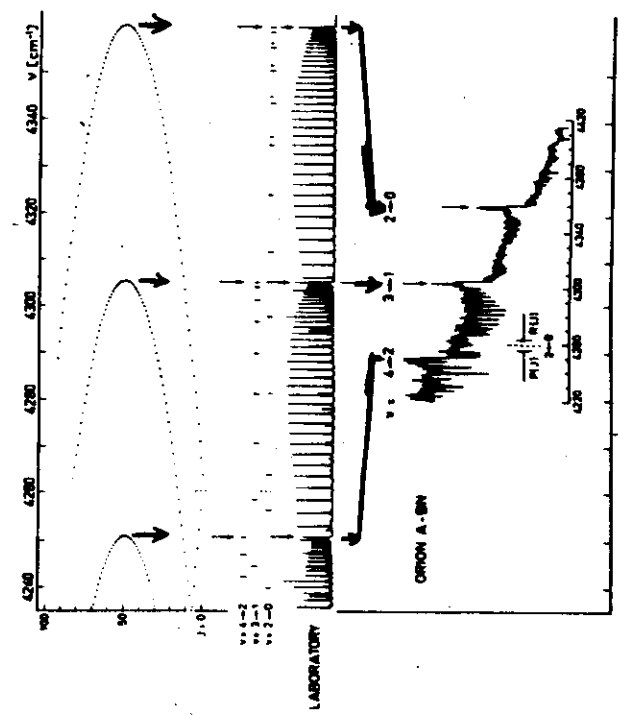
$$G_{01}(\text{CO}) \sim 2 \times 10^{-13} \text{ cm}^{-2}$$

$$R_{J, J-1} = \langle \sigma_{J, J-1} \rangle_{NH_3}$$

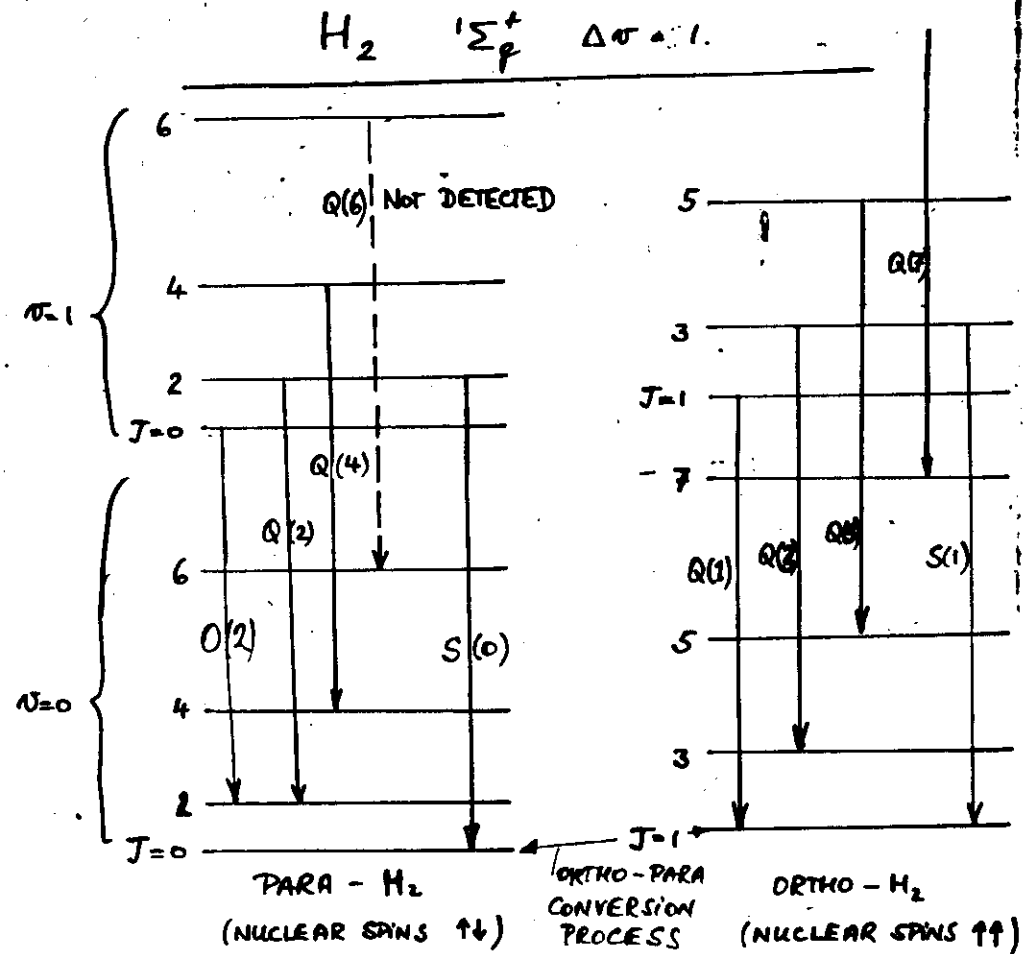
CO - H₂ 10 K

J	φ	1	2	3
φ	-	.393	.263	.011
1	.176	-	.193	.07
2	.276	.351	-	.09
3	.044	.00		

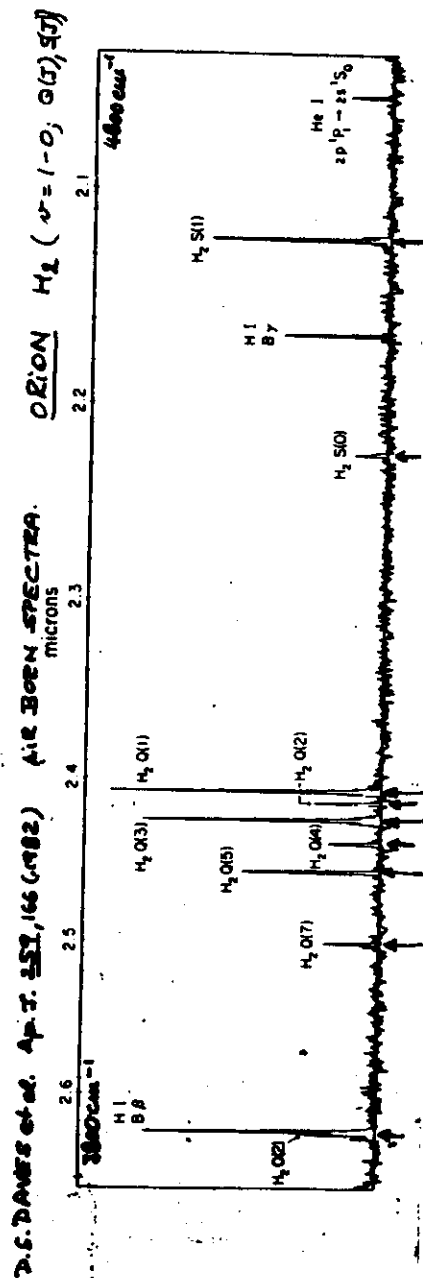
CO-OVERTONE BANDS



ENERGY LEVELS AND LOW J -TRANSITIONS OF

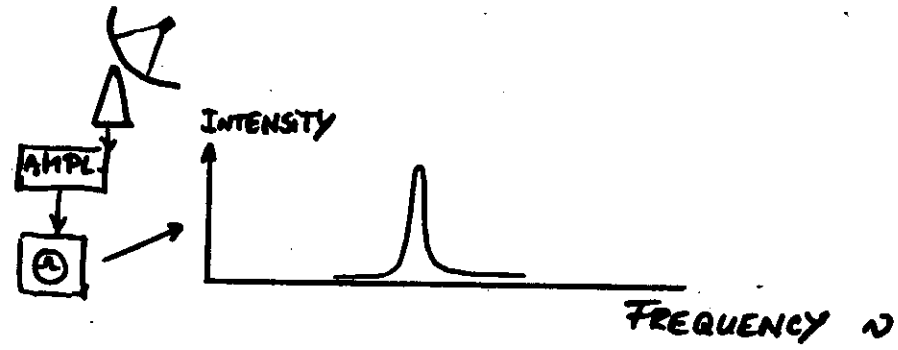
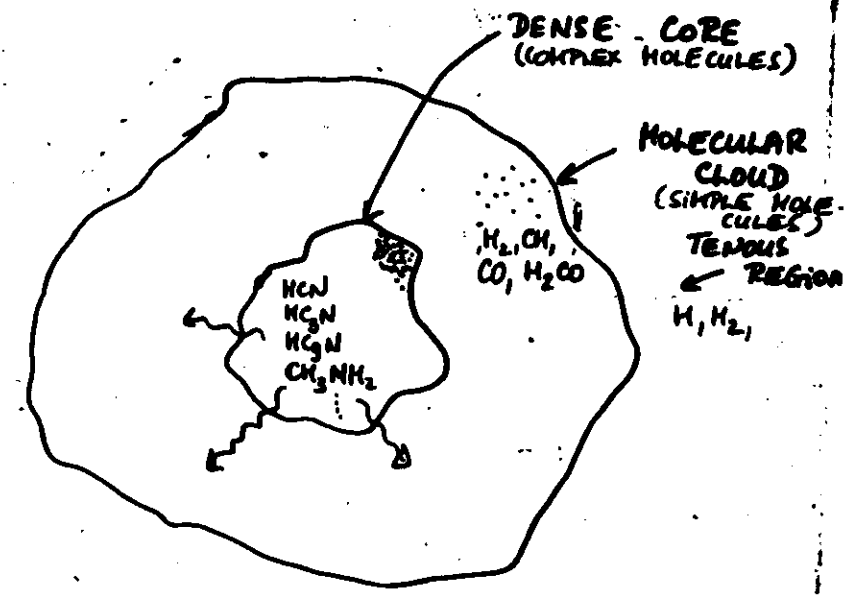


18



19

RADIATION FROM NEUTRAL CLOUDS MAINLY MOLECULAR TRANSITIONS

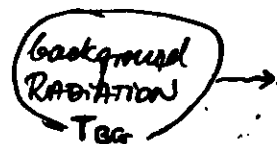


LINE BRIGHTNESS TEMPERATURE

$$T_B = \eta (J_\nu(T_{ex}) - J_\nu(T_{bg})) (1 - e^{-\tau})$$

η = beam filling
 τ = optical depth

ENERGY INPUT



ABSORPTION LINES

① $T_{bg} = 2.7K$

② RADIO CONTINUUM
(HI-REGIONS)

③ IR-CONTINUUM
(HOT DUST)

MOLECULAR LINE T_B

$T_{ex} < T_{bg}$: ABSORPTION LINES

① H_2CO , ^{13}CO , SiO
 $T_{ex} < 2.7K$
ANOMALOUS COOLING

② eg. NH_3 in DR21, Sgr B2

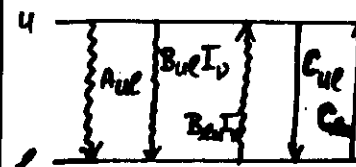
③ CO ABSORPTION IN ORION
 $N=2 \leftarrow 0; 1 \leftarrow 0$

④ Self Absorption
(MANY MOLECULES + RADIALS)

EXCITATION AND LINE FORMATION.

MOLECULAR GAS (H_2)
 T_{kin}
MOLECULAR TRANSITION
 T_{ex}

excitation temperature T_{ex}



COLLISIONAL EXCITATION RATE:

$$C_{ul} = n_{gas} \langle \sigma_{ul} v \rangle T_{kin}$$

RADIATION FIELD:

$$I_\nu = \langle I_{bg} e^{-\tau} \rangle + I_{rad}$$

$$n_u (A_{ul} + B_{ul} I_\nu + C_{ul}) = n_l (B_{lu} I_\nu + C_{lu}) \quad (1)$$

$$C_{lu} = C_{ul} \left(\frac{g_u}{g_l} \exp \left(-\frac{h\nu}{kT_{kin}} \right) \right) \quad (2)$$

$$\frac{n_u}{n_l} = \frac{g_u}{g_l} \exp \left(-\frac{h\nu}{kT_{ex}} \right) \quad (3)$$

LIMITING CASES

$$T_{ex} \sim T_{bg} \quad (I \ll 1; C \ll A)$$

$$T_{ex} \sim T_{kin} \quad (C \gg A)$$

$$J_\nu(T) = \frac{h\nu}{4\pi} \left\{ \exp \left(-\frac{h\nu}{kT} \right) \right\}$$

$$T_{ex} > T_{bg}$$

EMISSION LINES

(22)

FROM GAS AT $T \sim 10 - 3000$ K

EXCITATION CAUSED BY

GENERAL AND LOCAL ENERGY SOURCES

1. COLD CLOUDS (2.7 K, C.R., galactic interstellar background)

$$T_{kin} \sim 10$$
 K ; $n_{H_2} \sim 10^4$ cm⁻³

Example: Taurus cloudlets, ...

CO, NH₃, H₂CO, Radicals + IONS

(Note: possibly surrounded by warmer shells)

2. WARM CLOUDS: environment of embedded stars, or stars

$$T_{kin} \sim 15 - 50$$
 K ; $n_{H_2} \sim 10^5 - 10^7$ cm⁻³
(cloud close to star ~ 1 pc)

Example: W51, S140 (LTE-excitation)
W21, S106 (non-LTE-excitation)

CS, NH₃, HCN, (RADICALS AND IONS)

3. HOT CENTERS: environment very close to embedded star ~ 0.1 pc

$$T_{kin} \sim 100 - 600$$
 K ; $n_{H_2} \sim 10^7$ cm⁻³ - 10^8

Example: NH₃ in Orion (~ 200 K)

... CO high velocity wings (Orion, S140, ...)

... CO vibrational/high rotational transitions in Orion
 ~ 600 K.

(23)

4. MOLECULAR SHOCK FRONTS:

in regions of active star formation.

$$T_{kin} \sim 1000$$
 K - 3000 K

Examples:

... H₂ rotation-vibration line emission (~ 1500 K)

... CO Driftone Bandhead emission (~ 2500 K)

... NH₃ (J,K) > (9,9)

Note:

The HOT CENTERS and MOLECULAR SHOCK FRONTS are associated with high velocity gas flows.

5. CIRCUMSTELLAR SOURCES

HATS - outflow from evolved stars

$$T_{kin} \sim 100 - 1000$$
 K

$$n_{H_2} \sim 10^4 - 10^6$$
 cm⁻³

CO, HCN, ... RADICALS AND IONS

ion of the interstellar H_2O
 5_{23} has indicated a maser
 me molecular process must
 sively or to depopulate the
 a.

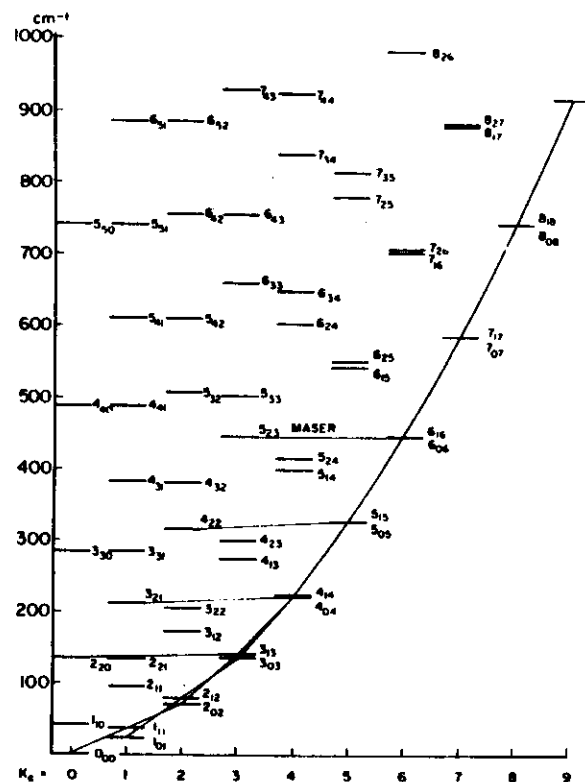
LEVELS

) that either the 6_{16} or the
 all other levels; there seems
 they happened to be very
 uate excessively a group of
 A levels) or depopulate a
 23 level. We divide many
 Group-A levels which are
 t levels which are composed
 levels with $K_a = J$ in which
 the C-axis. The justification
 by spontaneous emission is
 of H_2O sorted according to
 the bottom of each column
 hat spontaneous emissions
 ting from the $J_{0,J}$ and $J_{1,J}$
 ively, which also belong to
 the weaker transitions with
 fore if once molecules are
 energy the only radiative
 3 transitions. In particular
 is the first way out of the
 es Group-B levels there will
 increased induced emission
 roup-B levels via the maser

3 levels must be one of the
 re pumping and relaxation
 ciation process. Normally
 ional anomalies because it
 : for example, Townes and
 we wish to examine the

ROTATIONAL LEVELS

259



1. Rotational energy levels of H_2O molecules sorted out with K_c . The Group-A levels are composed of $J_{0,J}$ and $J_{1,J}$ levels which lie at the bottom of each column for a given K_c . The lines connecting various levels indicate spontaneous emission of the H_2O molecules. It should be noted that if molecules in Group-A levels relax from high energy levels the maser transition $6_{16} \rightarrow 5_{23}$ is the first connection between the Group-A and Group-B levels. Therefore if some molecular process depletes molecules in the Group-B levels there will be an inverted population between the 6_{16} and the 5_{23} levels.

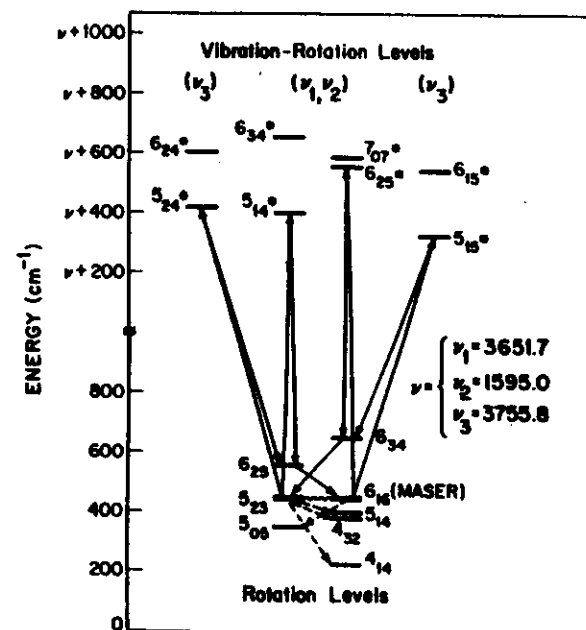
gy, as $4\Sigma^+$, $2\Sigma^-$ and 4Π .
 mon (1968), makes the
 peratures much greater
 J and H atoms close to
 curve. The switching
 rms $X^2\Pi$ OH. Second,
 requires about 1000°K,
 the pre-association rate
 association (the inverse
 don and Kopp (1971),
 $4\Sigma^-$ states. This rate is
 er A states. These rates
 ping, as seen in Figure
 rable to the maximum

vapor maser. Collisions
 to about $J=5$ if the
 re is likely to be about
 in either of the maser
 out 10^{-4} or 10^{-5} times
 n A-coefficients and
 igher rotational levels,
 temperature to about
 noted that the 6_{16}
 23 level. The observed
 itical depths. Because
 depths for the various
 large. The resonance
 appears to make the
 of 5_{23} (lower maser
 quate to pump these
 cause collisions across
 lizing effect. Infrared
 asterisks leads to
 1, making population
 vel can be shown to

naldehyde that was
 e-strength shown in
 d 1_{10} . However, the
 1_{00}^+ level does not

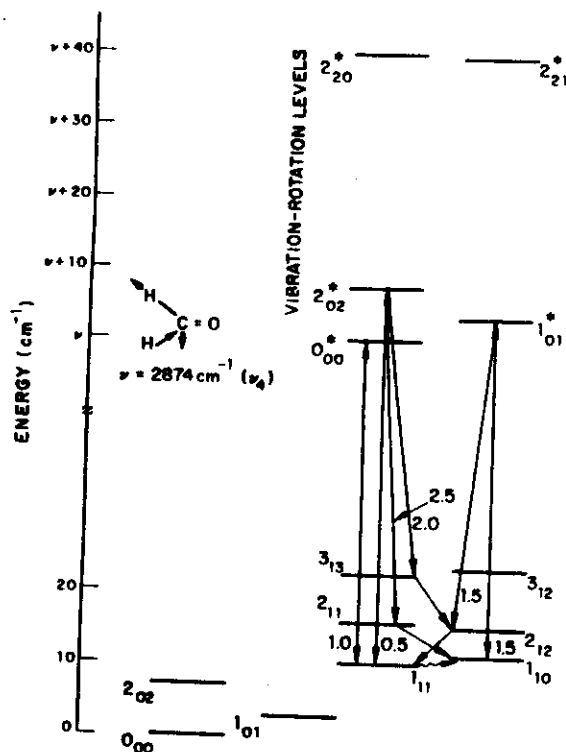
MASERS AND OPTICAL PUMPING

279



INFRARED PUMPING OF 22.2GHz H_2O MASER

7. Rotation and vibration-rotation levels involved in infrared pumping of H_2O maser (22.2 GHz). The lower rotational levels (many not shown) are in nearly-thermal equilibrium because of collisions.



8. Lower rotation and vibration-rotation levels involved in infrared pumping of formaldehyde (ortho) anti-maser at 6-, 2- and 1-cm wavelengths ($J = 1, 2$ and 3 doublets, respectively).

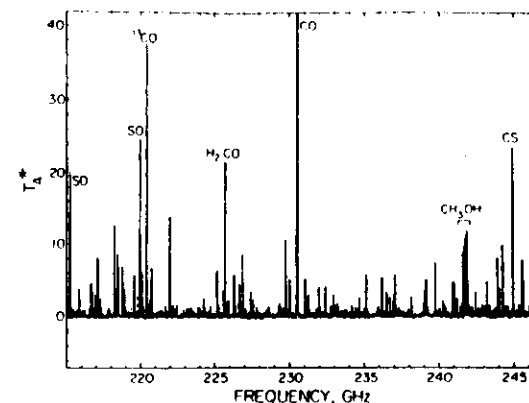


FIG. 1.—Spectrum of Orion A from 215 to 247 GHz. Data were obtained with a typical receiver noise temperature of 500 K (SSB) and backend coverage of 527 MHz. The original spectral resolution was 1.03 MHz (1.3 km s^{-1}). Spectral resolution in this plot is degraded by about a factor of 10.

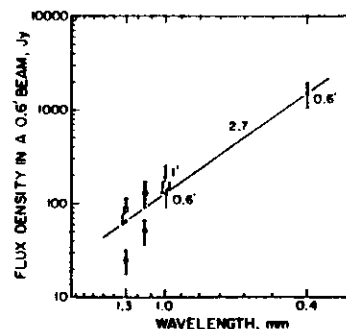


FIG. 2.—Broad-band and spectral line flux densities in a 0.6 beam centered on BN. Open circles show the line fluxes measured by Sutton *et al.* (1984) and Blake *et al.* (1984). Open boxes indicate the total broad-band fluxes measured simultaneously. Crosses represent the broad-band fluxes of Keene, Hildebrand, and Whitcomb (1982) and Elias *et al.* (1978), the latter both for a 1' beam and corrected for a 0.6 beam. Plus sign represents the extrapolation of these short wavelength measurements to 1.3 mm.

However, some additional uncertainties were introduced since the spectral line system was not well optimized for continuum observations. The measured broad-band flux was 96 Jy at 1.3 mm with an uncertainty of about $\pm 15\%$ on top of an overall calibration error of $\pm 20\%$.

This value for the broad-band flux can be compared with the various values in the published literature. The 400 μm emission from the core of Orion A has been measured and mapped with 0.6 resolution by Keene, Hildebrand, and Whitcomb (1982). They show that the submillimeter emission consists of an extended component plus two major compact

sources. The flux density at BN is 1500 Jy within a 0.6 beam, with an overall calibration uncertainty of $\pm 30\%$. Previously, the 1 mm emission from Orion was measured by Westbrook *et al.* (1976) and recalibrated by Elias *et al.* (1978). They derived a flux density of 188 Jy in a 1' beam. Assuming that the 1 mm emission has the same spatial distribution as the 400 μm emission, it is possible to estimate the 1 mm flux density in a 0.6 beam by using the mapping data of Keene, Hildebrand, and Whitcomb (1982). The result is an estimated flux density of 128 Jy for the smaller beam. Combined with the 400 μm result, this implies a spectral index of 2.7 between 0.4 mm and 1 mm. This is similar to but smaller than the spectral index estimated by Elias *et al.* (1978). Extrapolating with this spectral index to 1.3 mm, the wavelength of the spectral line observations, gives a flux density of 63 Jy in a 0.6 beam. The various broad-band fluxes are plotted in Figure 2. Comparison with the line flux indicates that 30%–40% of the total flux may be accounted for as line emission. This, however, is a lower limit since the line flux is based only on those lines stronger than 0.2 K.

III. DISCUSSION

a) Extrapolation to Weaker Lines

The calculations above have considered only those lines strong enough to be individually detected, i.e., those with peak antenna temperatures stronger than about 0.2 K. There will be a large number of weaker lines present which cannot be seen because of the noise level in the spectrum and because of the overlapping of stronger lines. Even at the 0.2 K level there are large regions of the spectrum in which the baseline cannot be clearly seen due to the overlapping wings of strong lines.

Some weaker lines can be predicted with some precision because they involve transitions with weaker matrix elements or are between more highly excited states of well-known species. But the general problem of predicting the number of

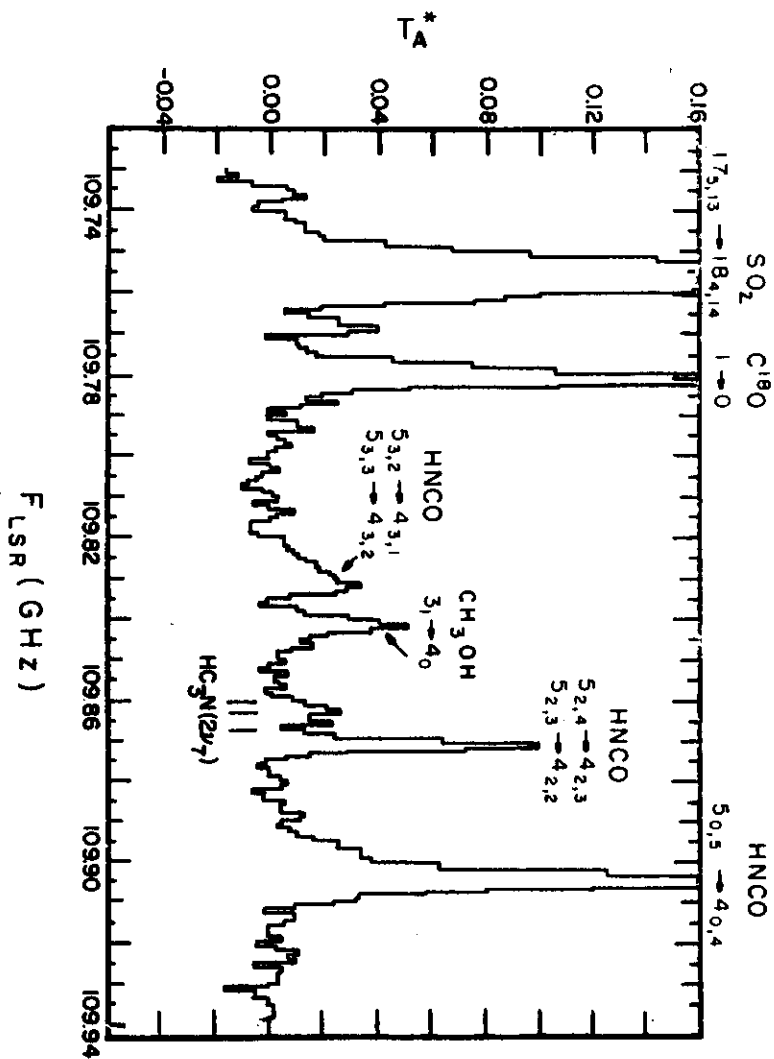


FIGURE 1

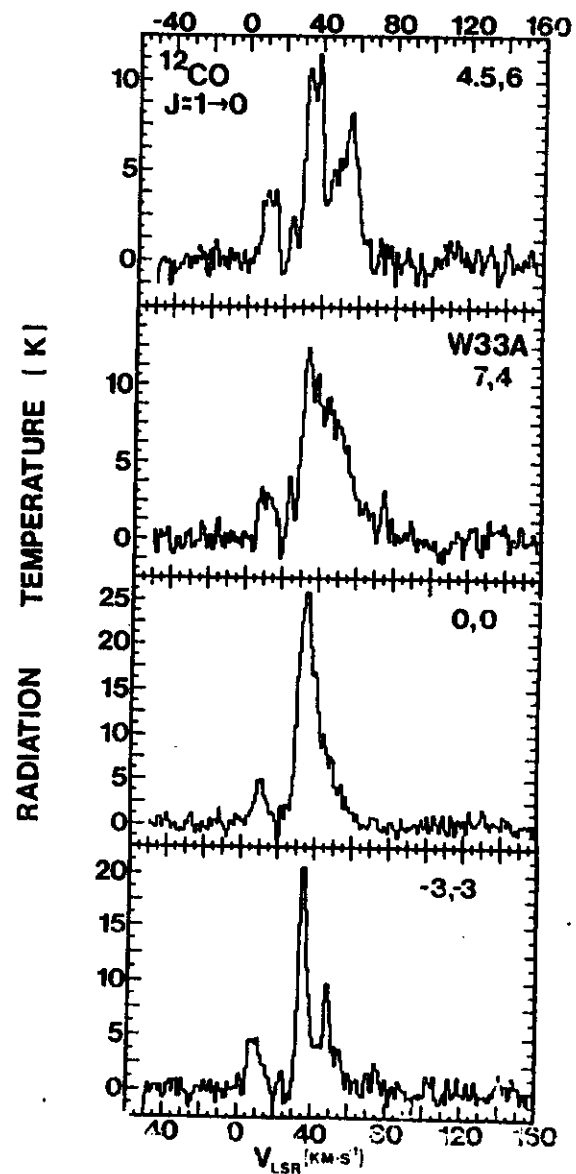


Figure 1

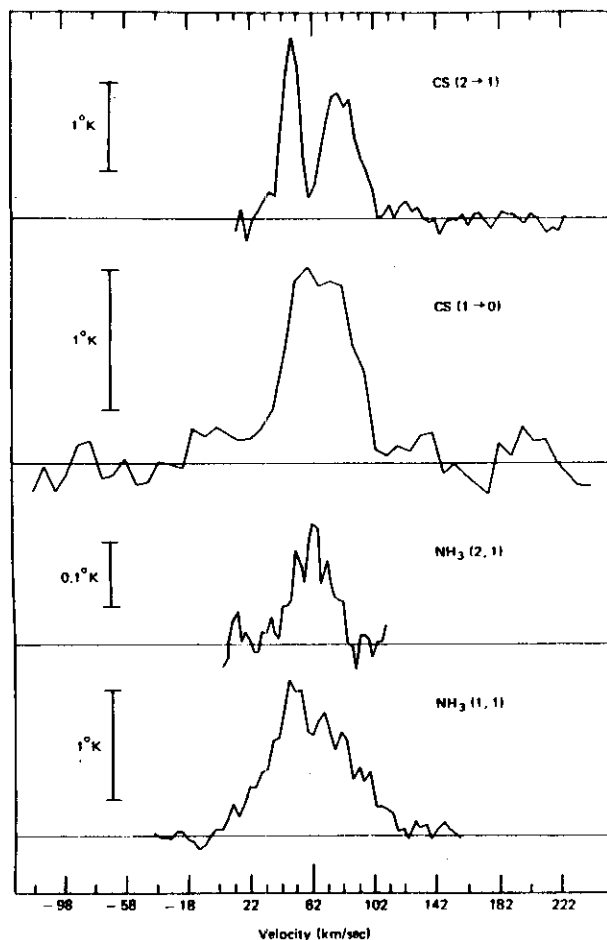


Figure 9.1 Spectra of NH_3 and CS in the Sgr B2 molecular source. The observation of the (2,1) line of NH_3 indicates that densities as high as 10^5 cm^{-3} may exist in the core of the source. The striking difference in the 2-1 and 1-0 lines of CS indicates that the Sgr B2 source may consist of two separate clouds or density condensations.

the line width. A secondary peak occurs in the HCO^+ line profile at a velocity of 5 km s^{-1} and at the velocity of the peak of the H^{13}CO^+ the HCO^+ line profile reaches a minimum at nearly the zero level. Even more severe self-absorption occurs near ρ Oph B2. In Figure 4, at a

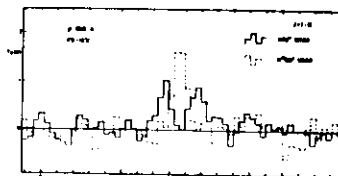


Figure 4. The $J=1-0 \text{ HCO}^+$ and H^{13}CO^+ line profiles toward a position $2'$ east of ρ Oph B2.

position $2'E$ of B2, the H^{13}CO^+ has twice the intensity of the HCO^+ and is centered between the two HCO^+ peaks. At the core of ρ Oph B2 we have obtained high resolution line profiles of the $J=1-0 \text{ HCO}^+$ and H^{13}CO^+ , $J=3-2 \text{ HCO}^+$ and $J=2-1$, $3-2$, $4-3 \text{ DCO}^+$ transitions (Fig 5). While different velocity resolutions were used for some of the lines, adjacent channels have been summed so that all of the final spectra have about the same resolution. All of the DCO^+ lines occur at 4.1 km s^{-1} and show no evidence of additional components. The HCO^+ lines, in contrast, show considerable structure. Both the $J=1-0$ and $J=3-2 \text{ HCO}^+$ profiles have absorption dips at slightly less than 4 km s^{-1} . The $J=1-0 \text{ H}^{13}\text{CO}^+$ also seems to have a very narrow deep dip which occurs at the velocity of the $J=3-2 \text{ HCO}^+$ peak. This is the only position which shows two component structure in the H^{13}CO^+ line profiles. All positions adjacent to ρ Oph B2 show only a single component, although at $2'N$ of B2 this component is shifted to a lower velocity, 2.9 km s^{-1} . Without further observations it can not be determined whether the H^{13}CO^+ is self-absorbed or two separate cloud components are present. The strength of the $J=3-2 \text{ HCO}^+$ and $J=1-0 \text{ H}^{13}\text{CO}^+$ at ρ Oph B2 relative to the $J=1-0 \text{ HCO}^+$ reveals that the latter is self-absorbed. The shift to a lower velocity also occurs in the DCO^+ lines at the position $2'N$ of ρ Oph B2.

5. PHYSICAL CONDITIONS WITHIN THE DENSE CORES

The observations reveal that there are a number of identifiable condensations within the ρ Oph cloud and that there is considerable variation in the physical conditions from region to region. In the $J=1-0 \text{ HCO}^+$ map several peaks, to the

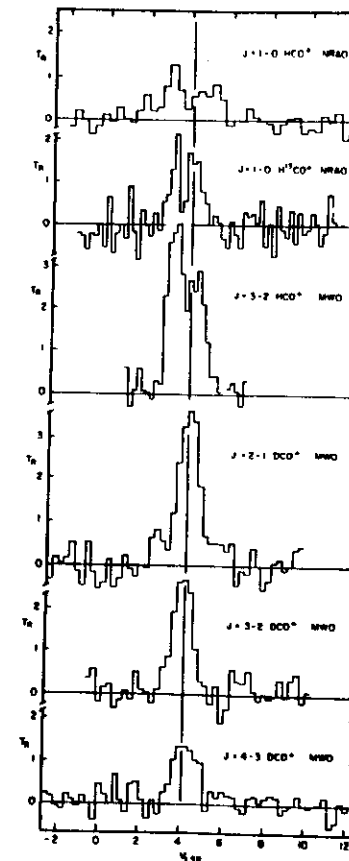


Figure 5. The line profiles of HCO^+ , H^{13}CO^+ and DCO^+ lines toward ρ Oph B2.

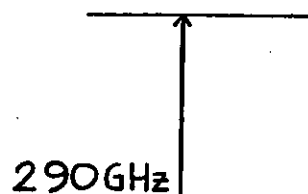
H₂CO energy levels (34)

$K_1=0$ (para)

$K_1=1$ (ortho)

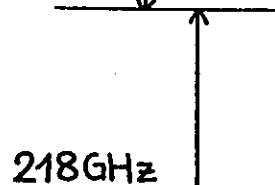
$J_{K_1 K_1}$

404



290 GHz

303



218 GHz

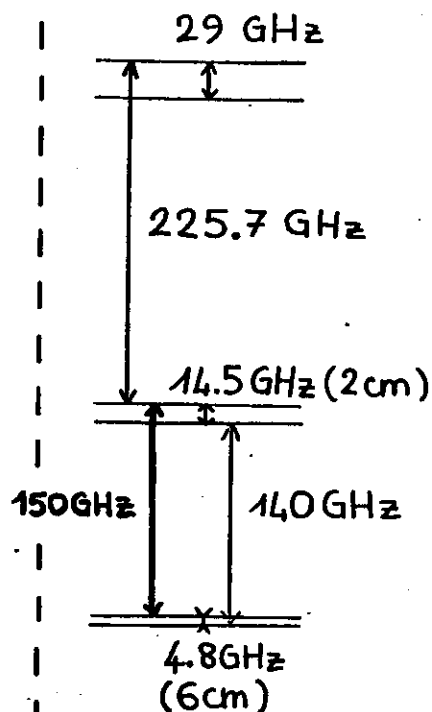
202

145 GHz
(2 mm)

101

72 GHz

000



29 GHz

225.7 GHz

14.5 GHz (2 cm)

150 GHz

140 GHz

4.8 GHz
(6 cm)

$J_{K_1 K_1}$

312

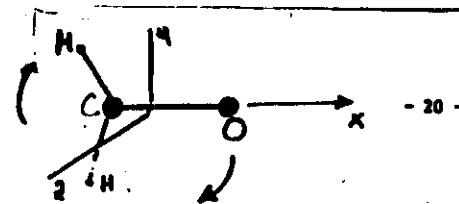
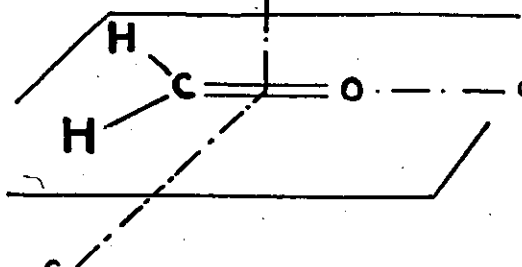
313

211

212

110

111



$J_{K_1 K_1}$

404

290 GHz
(1 mm)

303

218 GHz
(1.4 mm)

202

145 GHz
(2 mm)

101

000

PARA

H₂CO

$J_{K_1 K_1}$

312

313

29 GHz (1 cm)

226 GHz
(1.3 mm)

211

212

14.5 GHz (2 cm)

140 GHz (2 mm)

150 GHz (2 mm)

110

111

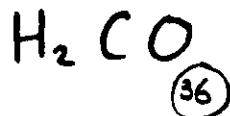
4.8 GHz (6 cm)

316 GHz

ORTHO

Figure B3 : Les premiers niveaux de rotation de la molécule H₂CO ainsi que les fréquences et les longueurs d'onde des transitions rotationnelles observées dans les nuages moléculaires. La flèche en pointillés indique la différence d'énergie entre le fondamental de l'orthoformaldéhyde et celui du paraformaldéhyde.

WHY MEASURE $[O]/[P]$?



$$[O]/[P] = 3 \times F(T)$$

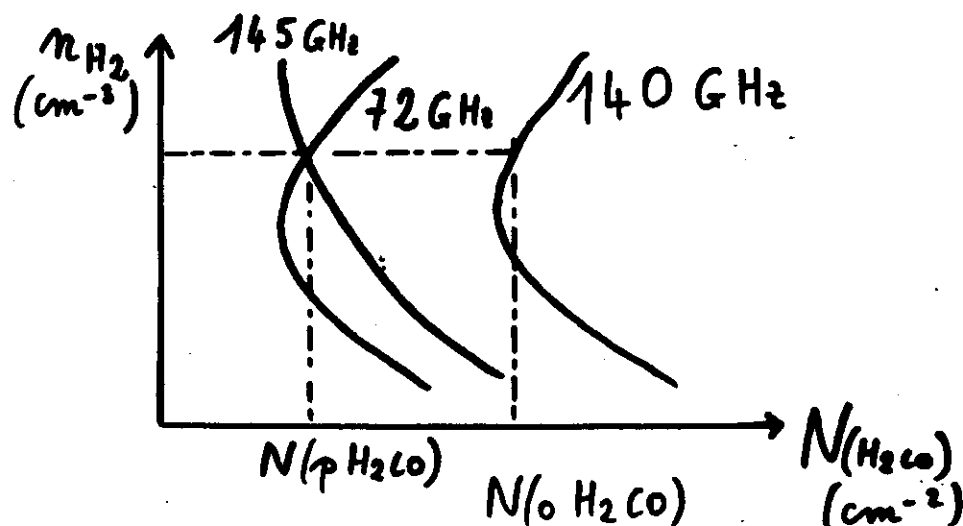
In The Laboratory $F(T) \equiv 1$

In the Interstellar Medium; $F(T)$ depends
of - H_2CO chemistry
- T of the clouds

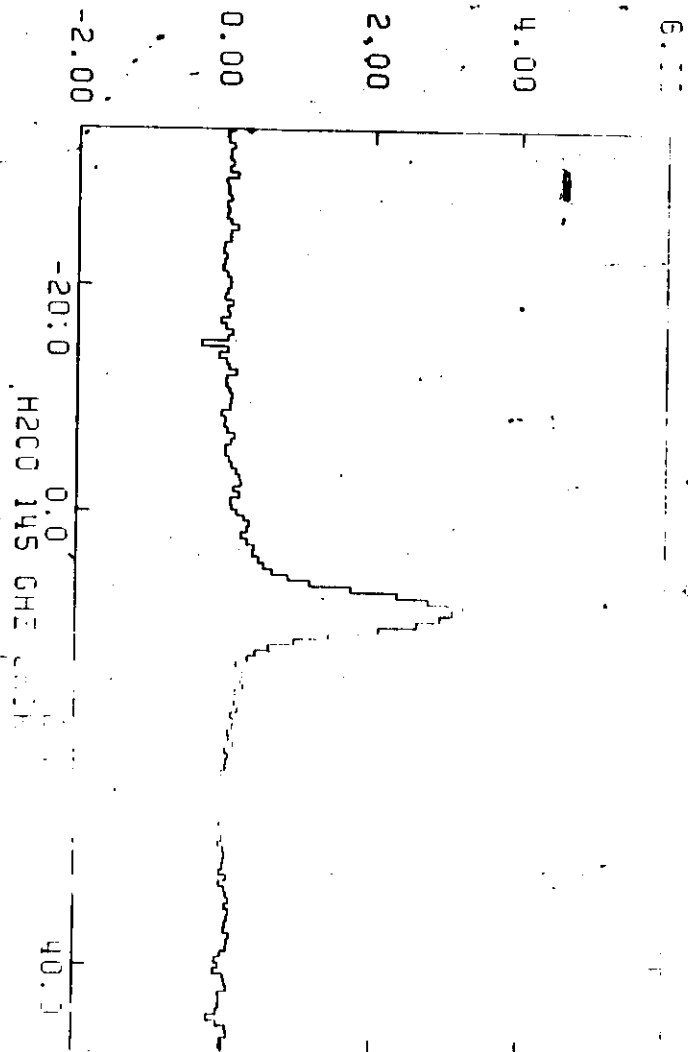
For every transition observed, plot of
 $T_B = f(n_{H_2}, N_{H_2CO})$. In Principle
you can measure (n_{H_2}, N_{H_2CO}) .

Pour chaque transition observée
on trace la courbe $T_B = f(n_{H_2}, N_{H_2CO})$

En principe, il y a une intersection
et on mesure (n_{H_2}, N_{H_2CO}) .



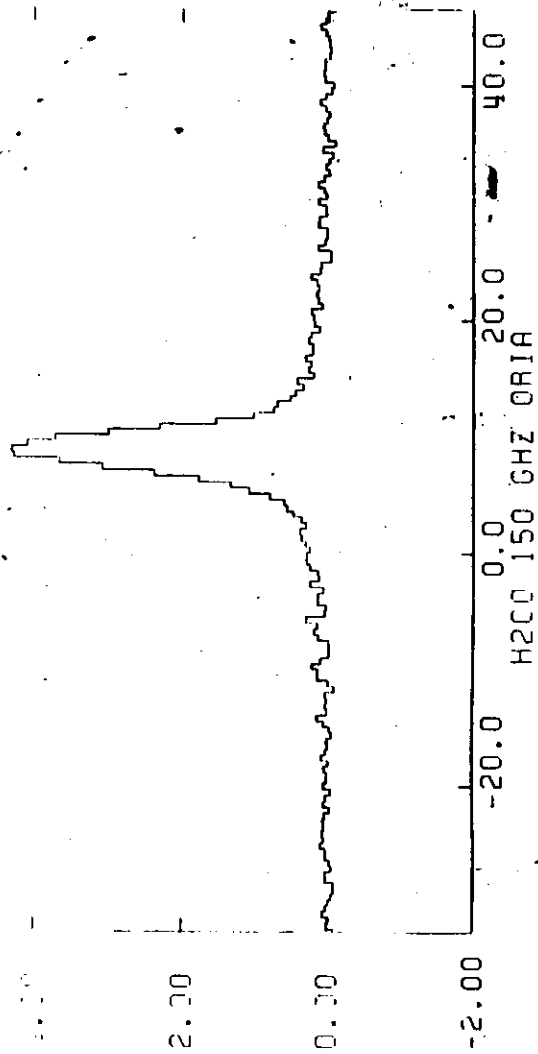
ANT TEMP DEGREES



DA. DEC (1968) -	5:32:47.8.	-5:24:29
L.B. (DECEMBER) -	208.965	-19.306
RAA CTRR (1968) -	5:32:47.8.	-5:24:29
RAA CTRR (JANUARY) -	"	"
SCANS USED	7662	7666
		7871

REST FREQUENCY = 148.602971 GHZ
CENTER VELOCITY (LSR) = 9.880 KM/S
CENTER CHANNEL = 129.59
REFERENCE CHANNEL = 3804.5
787 7889

.258 MHZ FILTERS
FREQ=Y
TIME = 3728.0 S



COPIES FILED
-204 MMZ FILTERS
FREQ
TIME - 6:00 PM '53

BEST FREQUENCY = 150.49369 MHZ
CENTER VELOCITY (LRR) = 9.898 KM/S
CENTER CHANNEL = 129.88
REFERENCE CHANNEL = 384.5
7845 7848 7851 7854 7857

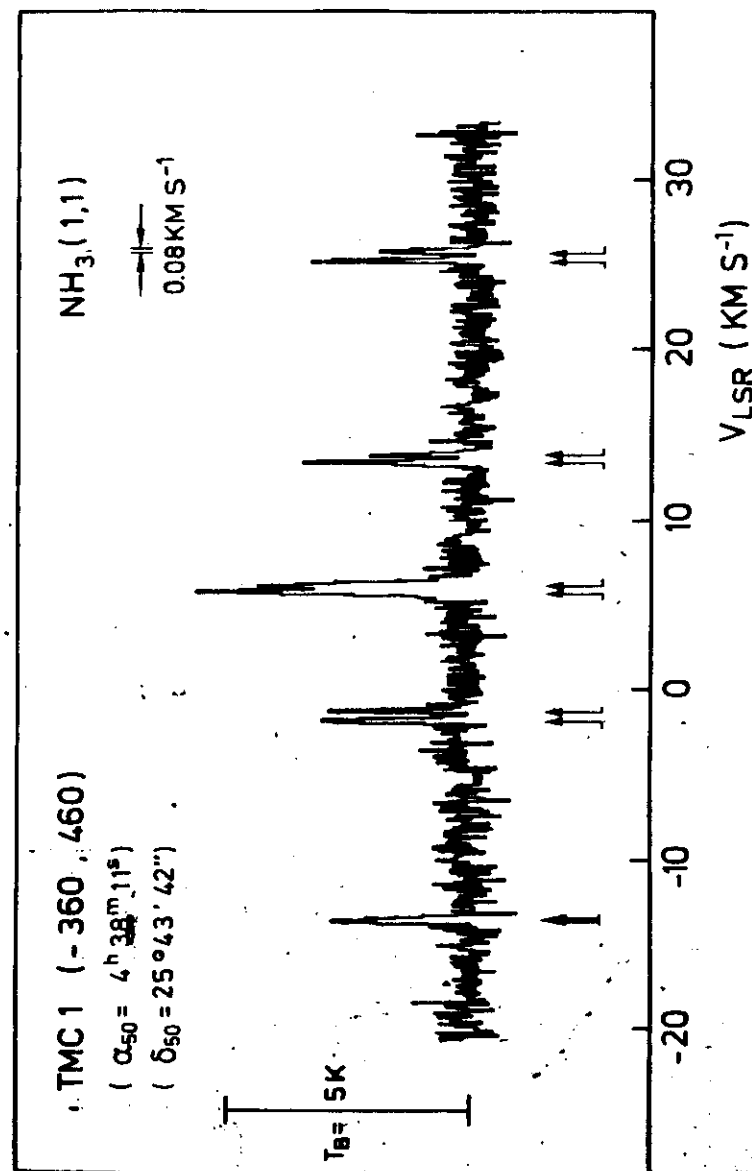
RA, DEC (1958) = 5 32 47.8 -5 24 29
L.S. (DEGREES) = 289.996 -19.386
MAP CTR (1958) = 5 32 47.8 -5 24 29
MAP OFFSETS (ARCMIN) = .0 .0
SCANS USED 7838 7833 7836 7838

DIFFICULTIES:

- optical depth of H_2CO 2 mm lines
- fragmentation of the cloud
- size of the beam at different frequencies
- is chemical equilibrium reached?

CONCLUSIONS:

- $[O]/[P] \sim 3$ in ORION
- $[O]/[P] \sim 1.5$ in Low Temperature clouds
- fragments needed in ORION
- isotopic ratio smaller in ORION ($^{12}C/^{13}C$)



c) The broadening of the line profile:

(42)

In molecular clouds, the linewidth is determined by Doppler-broadening:

- gaussian thermal broadening
- turbulent
- systematic motions

Only in the case of rotation, it appears that turbulence is the main cause of broadening, and that the line classes are suprathermal, but still gaussian.

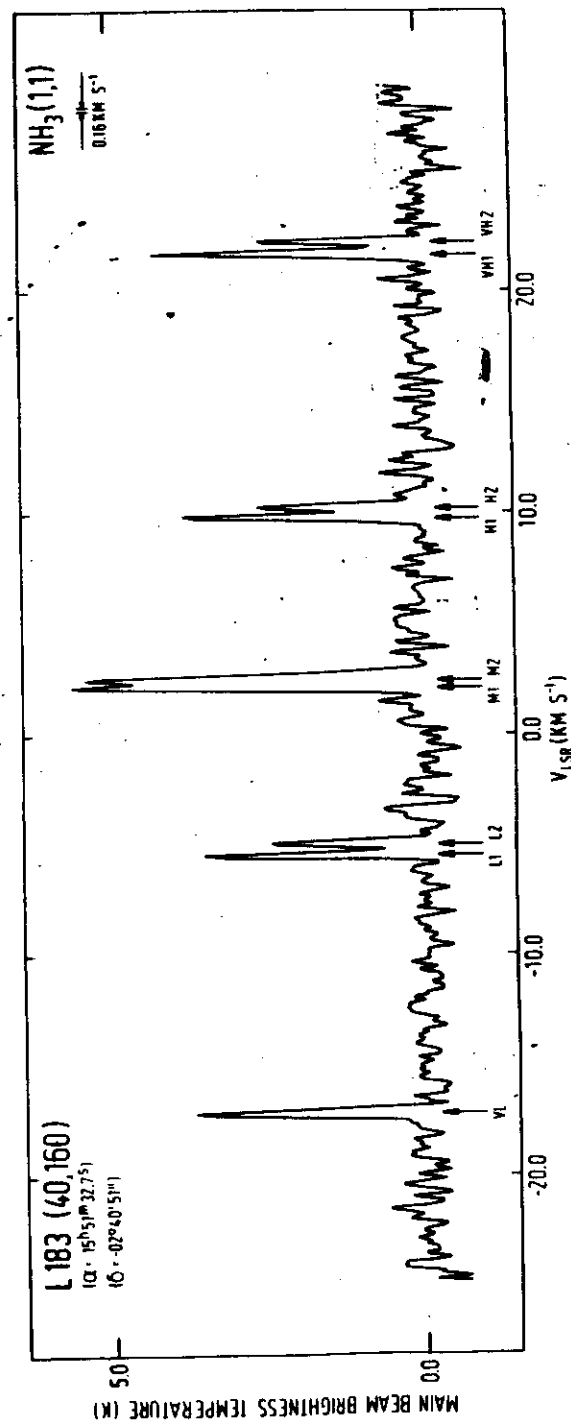
If you look in the same cloud two different transitions for two different molecules, you may be able to measure the kinetic temperature of the cloud T_k :

$$\begin{aligned} \text{In L 183} \quad \Delta v_1 &= 0.21 \text{ km/s for } \text{NH}_3 \\ \Delta v_2 &= 0.17 \text{ km/s for } \text{HC}_3\text{N} \end{aligned}$$

$$T_{\text{kin}} = \frac{\Delta v_1^2 - \Delta v_2^2}{8 \ln 2} \frac{\eta_1 \eta_2}{\eta_2 - \eta_1} = 9.4 \text{ K}$$

Another cause of broadening is the case of large optical depth (see Modern Aspects of Microwave Spectroscopy by CHANTREY for a detailed analysis).

CAUTION: reliability of this method may be questionable
See IRC + 10216 spectra of different molecules!!



(43)

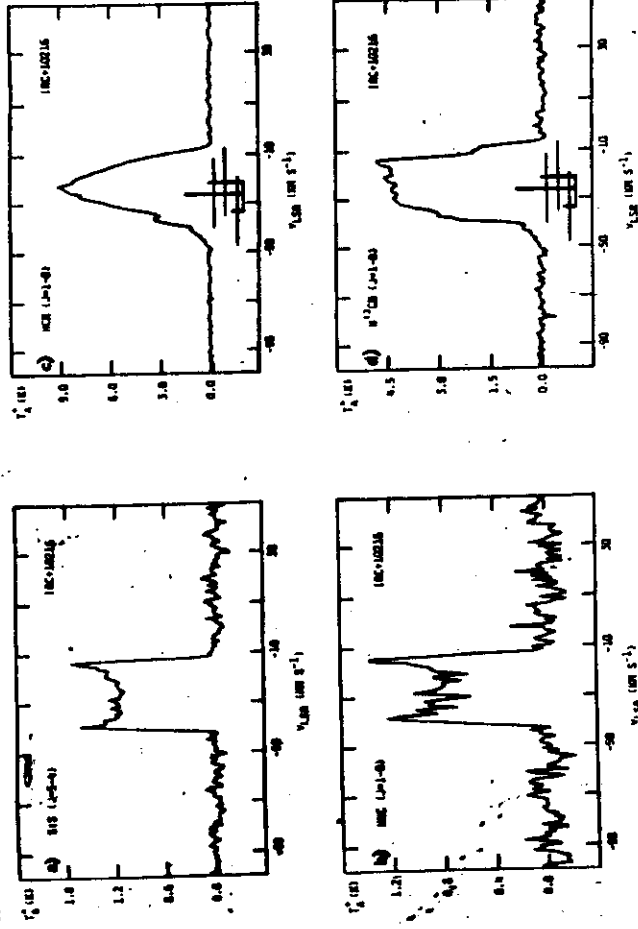


Fig. 4. a) SiS ($J=4-3$), b) SiO ($J=1-0$), c) HCN ($J=1-0$), and d) H¹³CN ($J=1-0$) spectra of IRC + 10216 (velocity resolution ~ 0.85 km s⁻¹). The center velocities, the full widths at zero power (determined from the scalar velocity and the gas expansion velocity derived in Sect. V), and the LTE ratios for the three quadrupole components of HCN and H¹³CN have been enlarged.

An upper limit (3 σ) may be placed on the SiO ($v=1$, $J=2-1$) (iv) HNC, $J=1-0$, Fig. 4b.

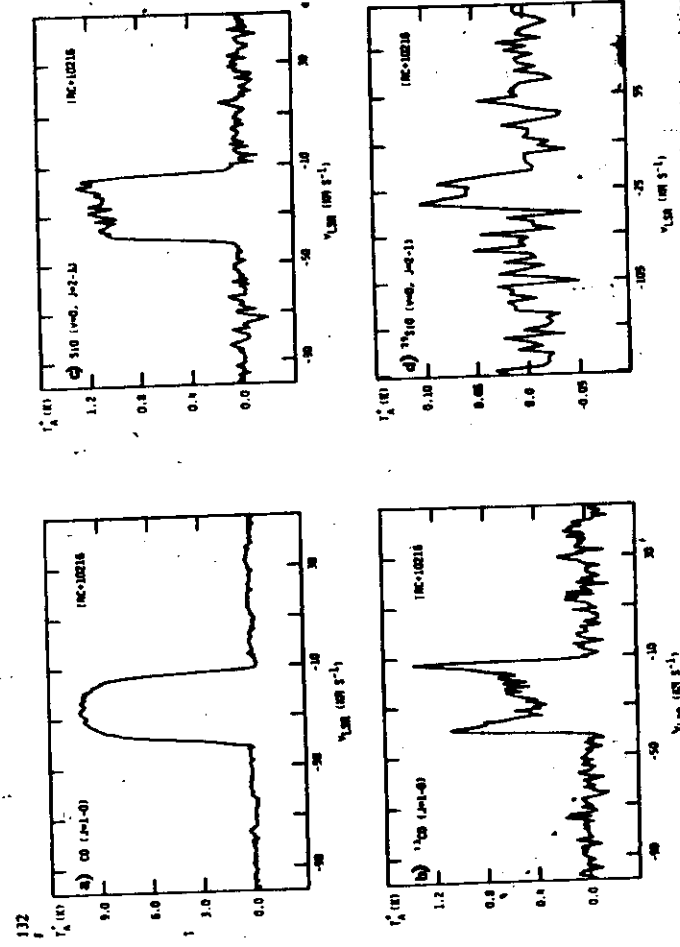


Fig. 3. a) CO ($J=1-0$), b) ¹³CO ($J=1-0$), c) SiO ($J=1-0$), and d) SiS ($J=4-3$) spectra of IRC + 10216 (velocity resolution ~ 0.67 km s⁻¹ in a and b, 0.86 km s⁻¹ in c, and 3.5 km s⁻¹ in d).

where

$$R_{\nu} = 2(\ln 2)^{1/2} R_{\nu}/\Delta\nu$$

We have in Fig. 2 also entered values for $T_{\text{MB}}(v=0, J=1-0)$

(i) CO and ¹³CO, $J=1-0$, Fig. 3a and b

The ¹³CO and CO profiles are excellent examples of spatially resolved optically thin and thick emissions. The antenna temperature of the ¹³CO doublet is about 3.

300-K

200-K

100-K

— l_{10}
— l_{11}
— l_{01}

— 0_{00}

— 2_{20}
— 2_{21}

— 2_{11}

— 2_{12}
— 2_{02}

— 3_{31} 3_{30}

— 3_{21}

— 3_{22}

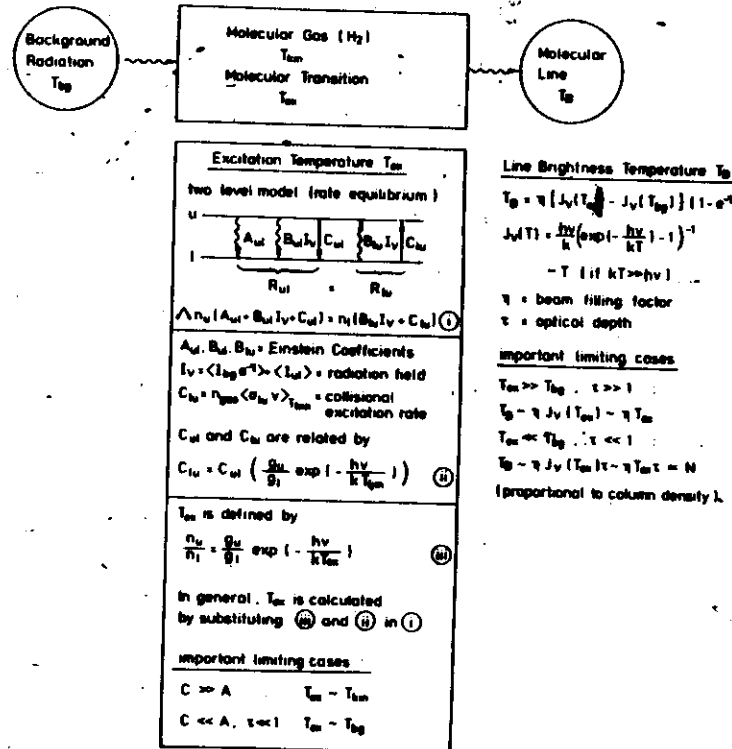
— 3_{12}

— 3_{03}

$H_2 D^+$

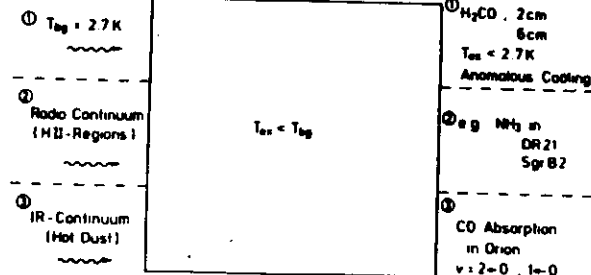
46

EXCITATION AND LINE FORMATION



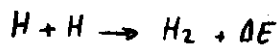
47

Absorption lines



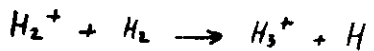
Interstellar chemistry:

The density is so small and the temperature so low that the chemistry of the interstellar medium has little to do with the laboratory chemistry. The binary collision between atoms, or between atoms and ions will therefore be the building block of the heavy molecules. But certain types of collision do necessitate a third body (dust grain for example) to recede the energy: this is the case for the most important one (molecular hydrogen).



DE must be absorbed by a 3rd body

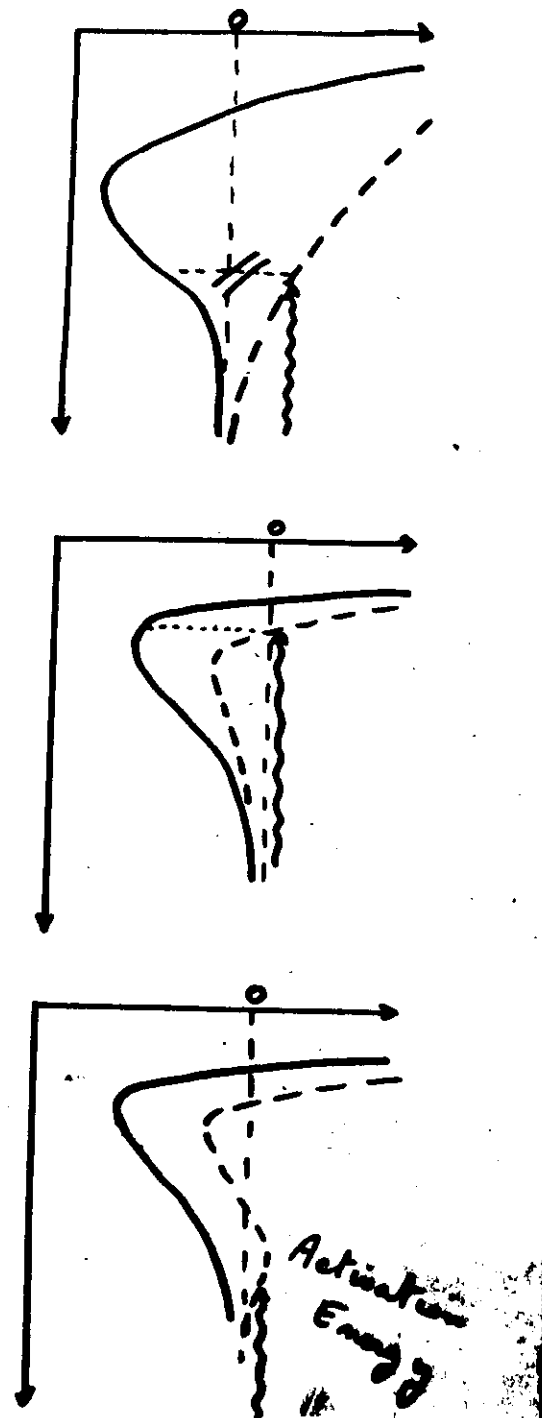
In the dense molecular clouds, exposed to the U.V. radiation field, the COSMIC RAYS do ionize the major constituents and produce H^+ , H_2^+ , He^+ which will react rapidly with CO , H_2 ... to build more complex molecules.

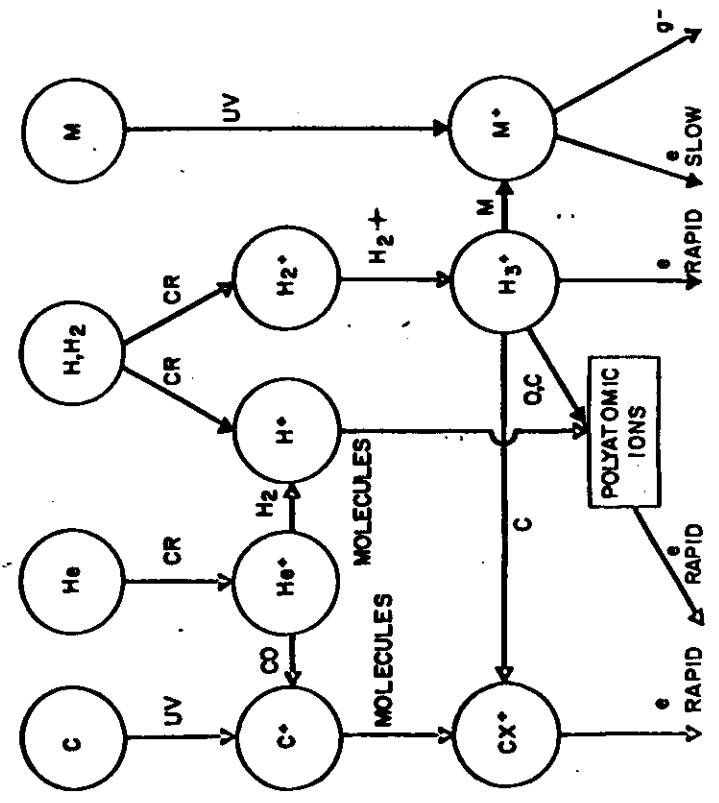
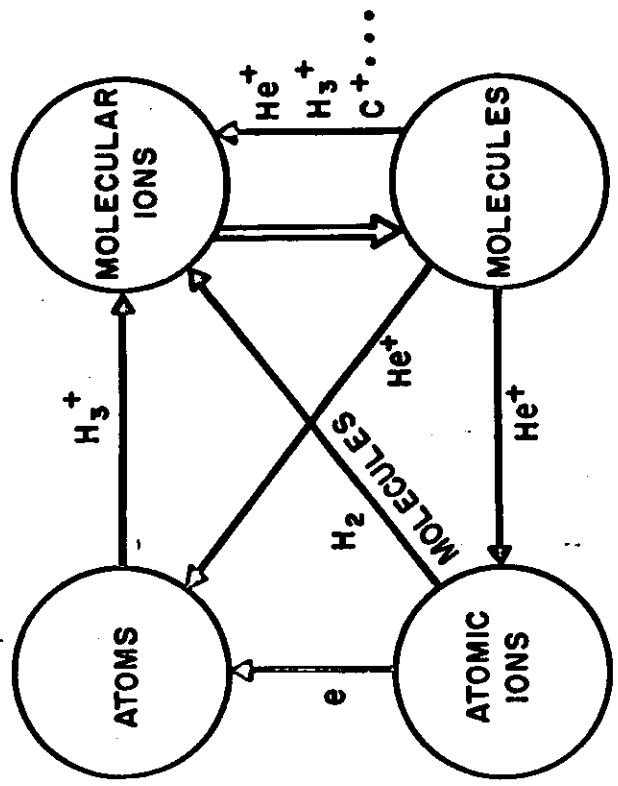


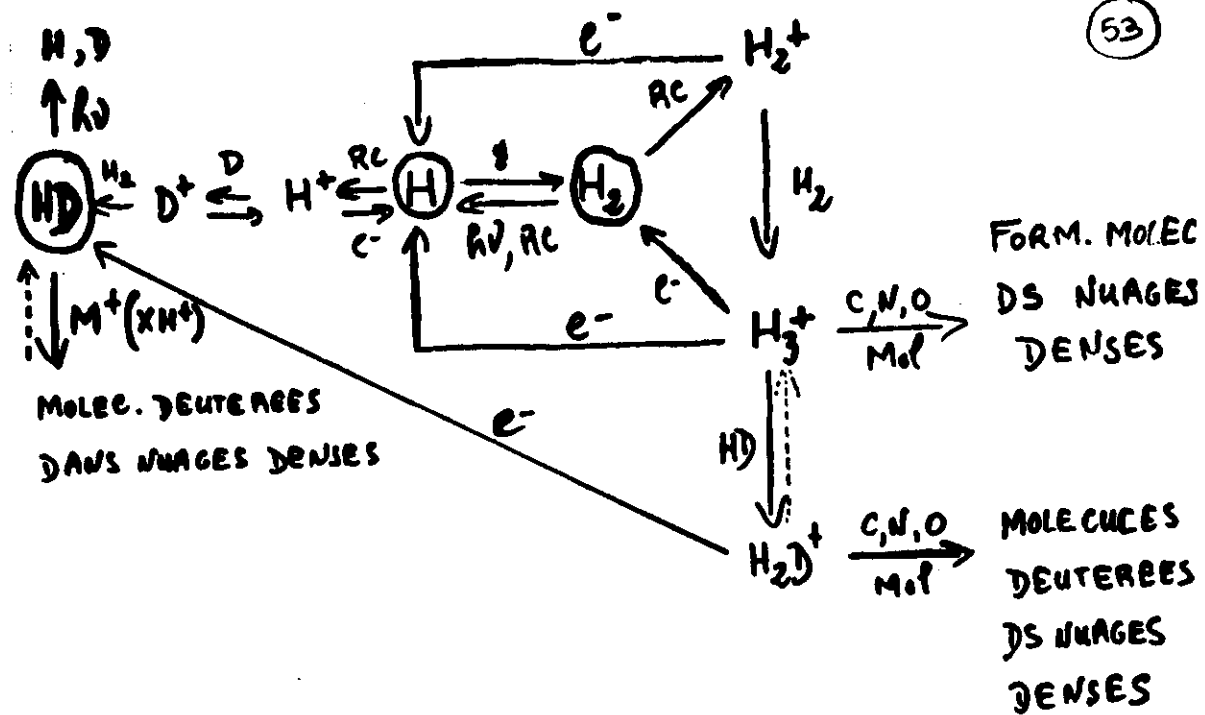
Some of the possible schemes are presented in the following transparencies. The models used for predicting the abundances of the light molecules are close to the measured abundances. However H_2 and H_2CO do not fit this scheme and must be fabricated on dust grains (physiosorption on the surface of the interstellar grains).

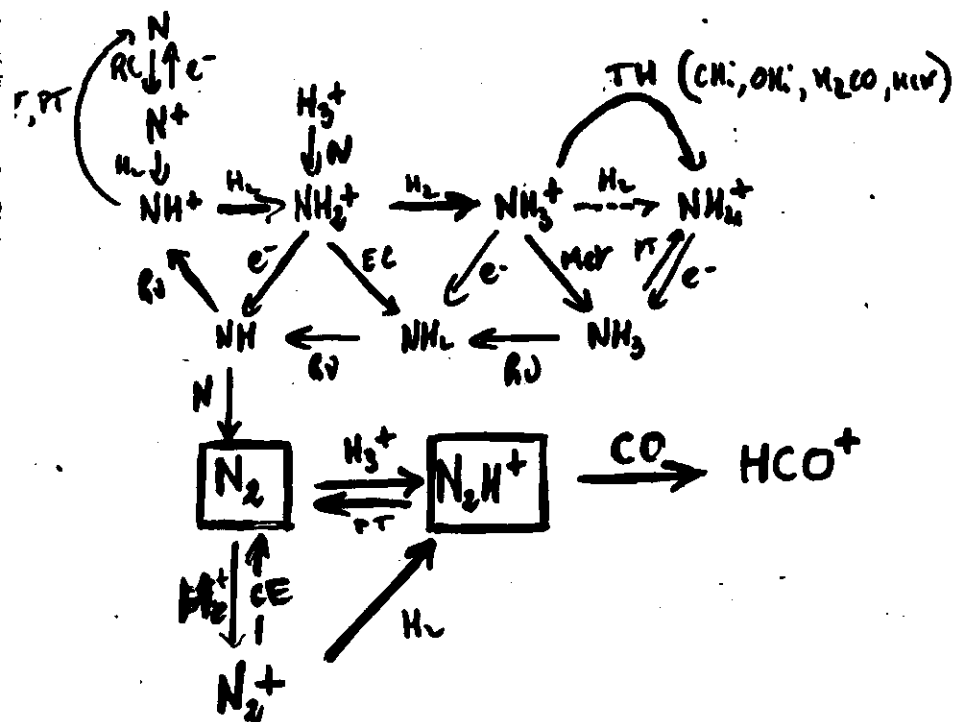
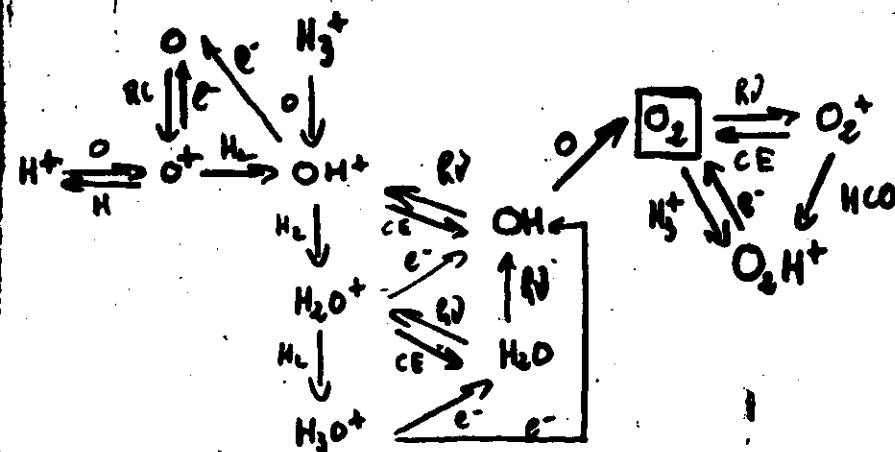
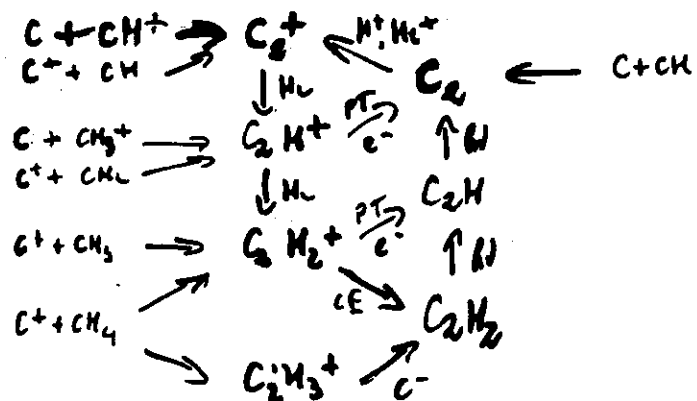
One of the major success of gas phase chemistry has been the prediction of the enrichment in deuterium of deuterated molecules: I will elaborate on DCO^+ and D_2CO respectively:

DISTANCE BETWEEN NUCLEI

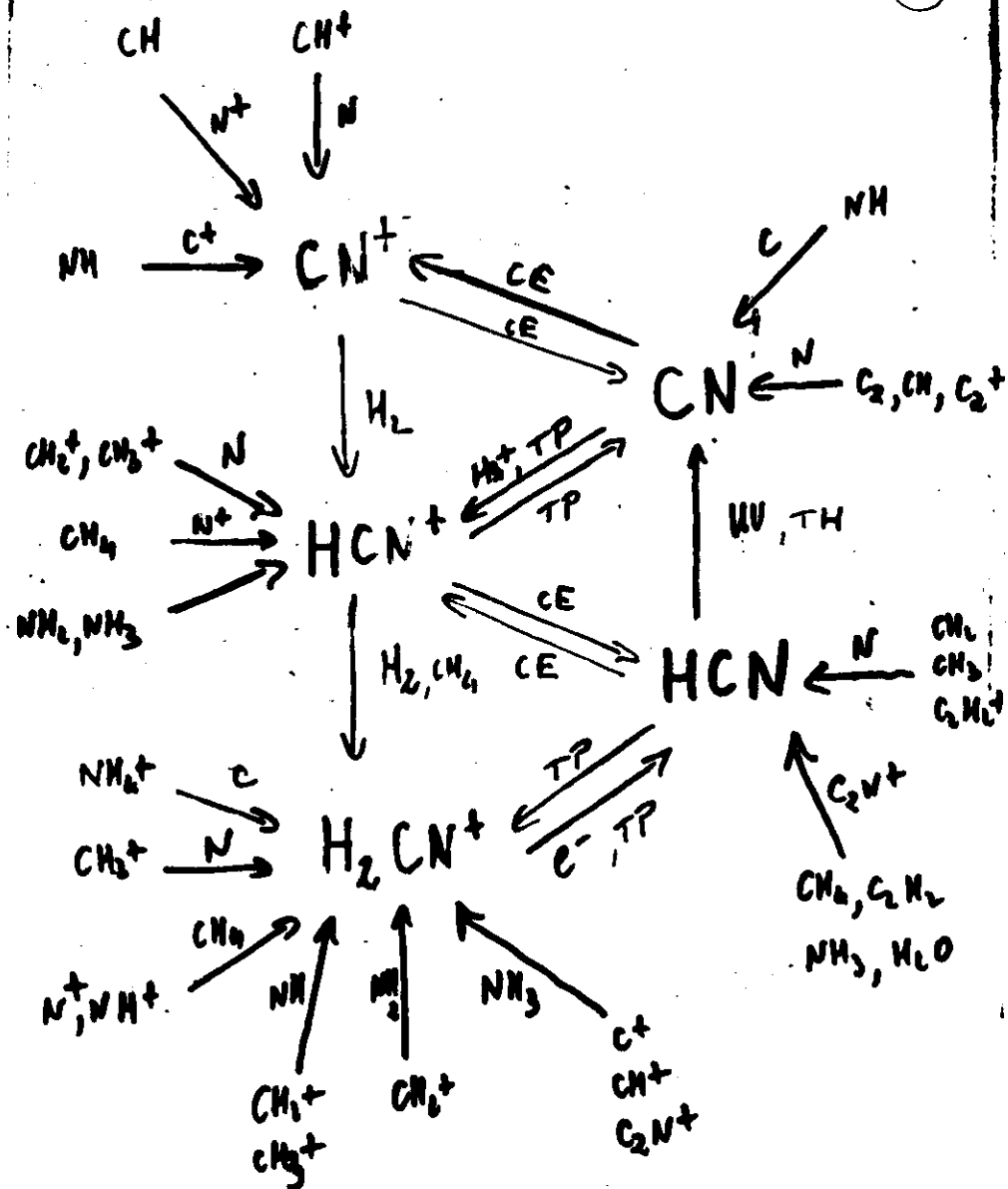




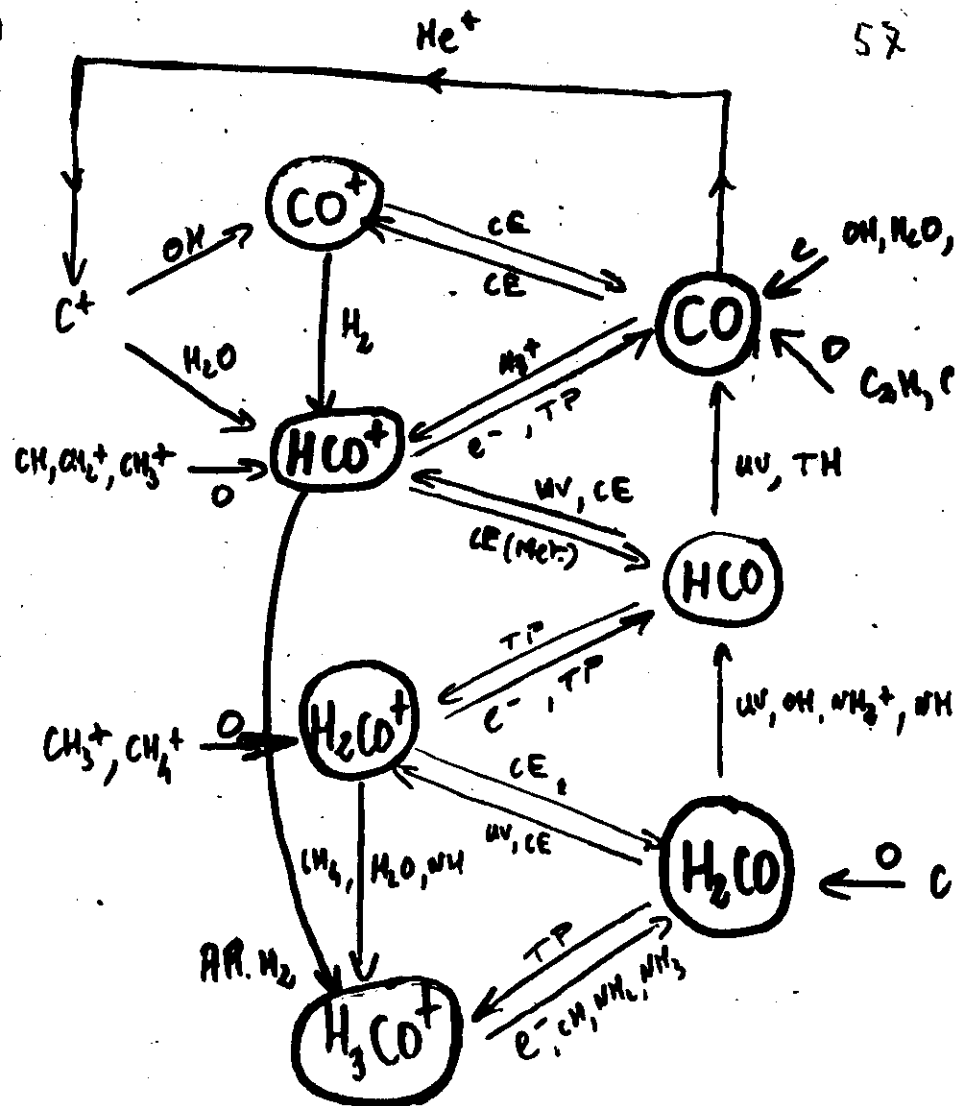




56



57



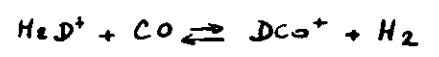
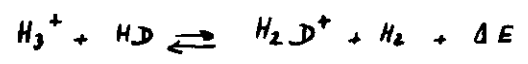
TH: Transfert d'un at. H

CE: échange de charge

TP: Transfert de Proton

UV: Photoionisation - Photodissociation

AR: Arrêt radiatif



DCO^+ should be enriched compared to HCO^+ , and this is the case (slide). This chain of reaction is controlled by the local density of electrons in the molecular cloud, and permit a precise measurement of the electron density in the cloud.

Watson has also shown that in the case of CO, the exchange reaction

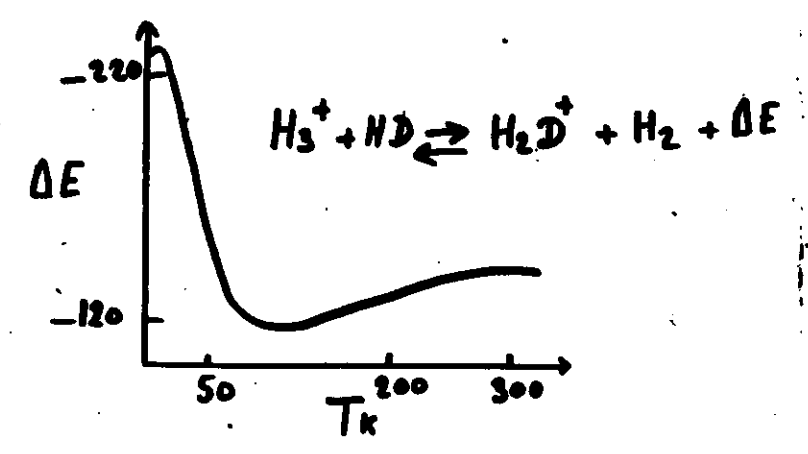


$$\Delta E \sim k \times 60 \text{ K}$$

explain the different CO/ ${}^{13}\text{CO}$ ratio observed in the center of molecular clouds and at the boundaries of such a cloud (in particular for $T_K \approx 10 \text{ K}$).

The recent detection of H_2D^+ (Philippe, 1985), the abundance of DCO^+ clearly indicate that gas phase chemistry is an important tool in the building of complicated molecules. It must be said that there are cases where we do NOT understand how to manufacture molecules: the but ions are CH and H_2CO which seem to be overabundant by 2 or 3 orders of magnitude!

The case of cyanopolyynes (HC_nN is observed) is also far from being solved.



D] A MAJOR DISCOVERY:

The behavior of a small grain (of ~ 50 atoms) being hit by a single U.V. photon led A. Leget and J.L. Pigot to suggest that grains of this size would rather be close to coronene ($C_{24}H_{12}$). In comparing their spectroscopic properties, they identified most of the Infrared (2-12 μm) unidentified features which had been known for >15 years. They also predicted that these polycyclic aromatic molecules could emit strongly in the 10 μm band of the IRAS satellite, and this has been recently confirmed (see La Reiche, March 1985).

Their next suggestion is that diffuse bands, which were discovered 70 years ago and which have remained a mystery until today are caused by these molecules. Many groups are presently working on this suggestion, and hope to have the answer in the coming year.

E] A SIMPLE CASE:

The OH radical has been discovered 10 years ago in the gaseous cloud associated with a comet by Gerard and coworkers. On his trajectory, a comet has a velocity relative to the sun which changes with time, and therefore the U.V. excitation of OH can be tested (the Fraunhofer spectrum of the sun is Doppler shifted). With a single model, it can be shown that period of emission and absorption of Λ doublet of OH should occur at specific velocities, and this has been precisely verified on more than 10 comets (slide).

LETTER

L22

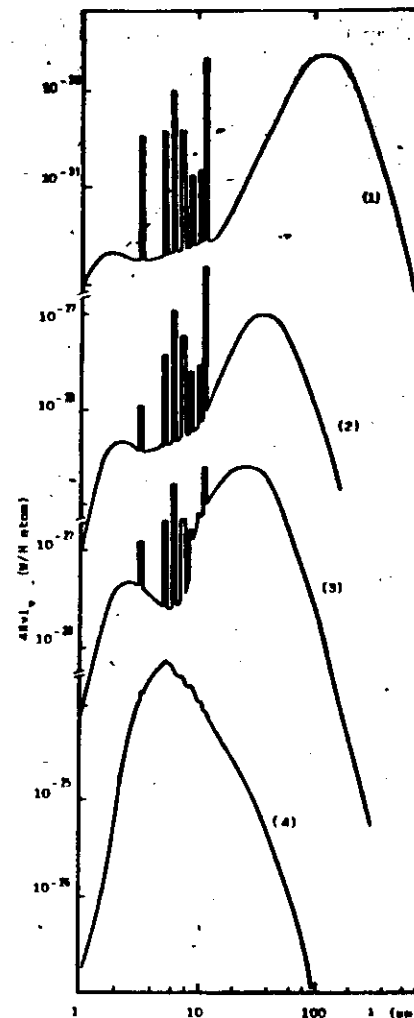


Fig.4 : Spectra obtained for different incident radiation fields :
Spectrum 1 : Local Interstellar Radiation Field
Spectrum 2 : 1pc from an O5 star
Spectrum 3 : 0.33 pc from an O5 star
Spectrum 4 : 100 stellar radii from a M0 star

This process should help to measure the rotation period of a comet on itself. (62)

What are the interstellar molecules used for

(63)

A) Star Formation: Physical parameters of molecular clouds

The Molecular Clouds are the places where stars do form after the cloud collapses on itself. It is therefore extremely important to measure the size of the fragments and their density. These objects being opaque to optical radiation can only be investigated with molecular spectroscopy.

The following Kampanian show the size of the fragments measured by H. Pencil and coworkers using a new technique, and their relation to the thermodynamics of the collapse. 3 sizes seems to fit best with the data.

The electron density n_e , related to the Cosmic Ray flux penetrating these clouds, is measured from the deuterated molecules and their exchange reaction rate. A value $\frac{n_e}{n_{H_2}}$ close to 10^{-8} is deduced by Guiché et al, and explains quite well the ionic chemistry going on.

The magnetic field can be measured directly by Zeeman effect on molecules like OH (27% ground state). After a decade of painstaking efforts, fields of the order of 11 to 28 μ Gauss are measured in a few galactic zones. If we recall that from Faraday rotation, the longitudinal magnetic field measured by pulsars is of the order of 3 μ Gauss, the enhancement in molecular clouds does

CAS A
HCO⁺, J=1-0

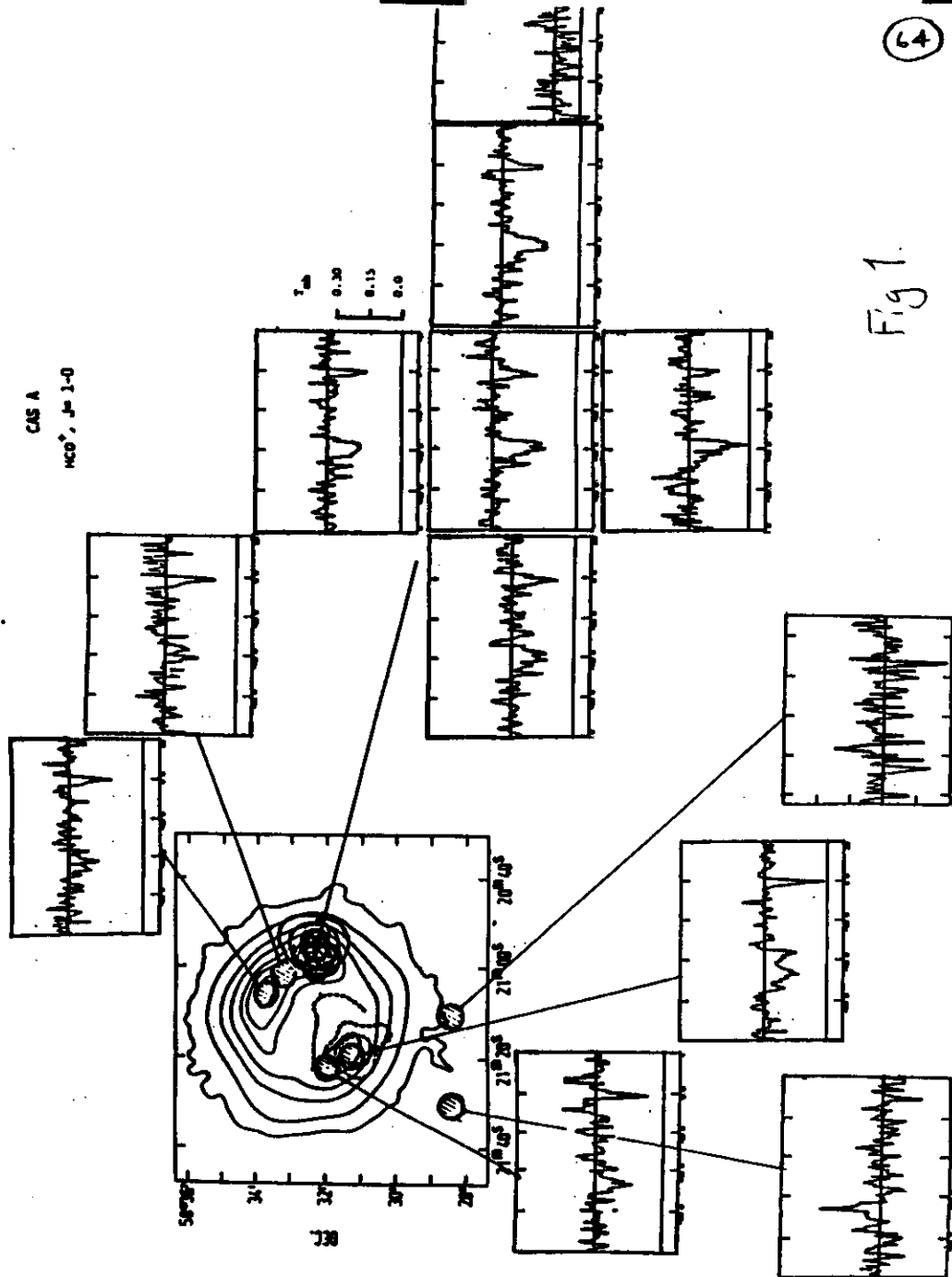


Fig 1

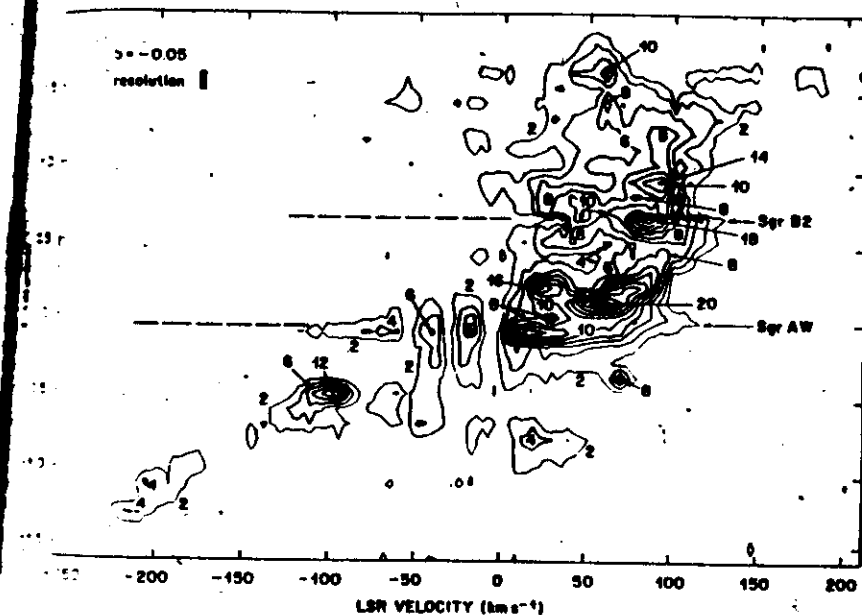


FIG 2. HCO⁺ at $b = -0.05^\circ$, $l = 2^\circ - 15^\circ$. Contours are equally spaced every $\Delta T_b = 0.2$ K and are labeled in units of $(T_b \text{ per } 0.1 \text{ K})$. Note the emission at -51 , -38 , -2 , and (perhaps) 32 km s^{-1} . The dashed lines at the positions of Sgr B2 and Sgr AW indicate the velocities at which the Orion and Perseus arms absorb from the continuum. Presumably low-excitation HCO⁺ is present in the region between these dashed lines and extends to other galactic longitudes as well.

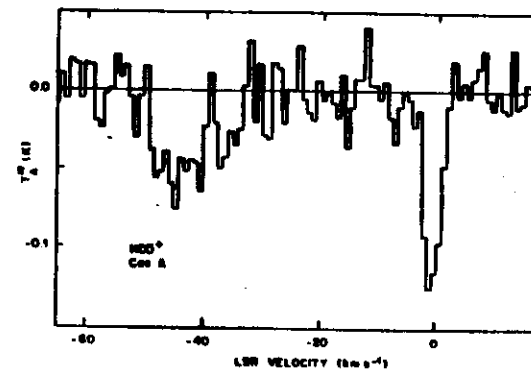


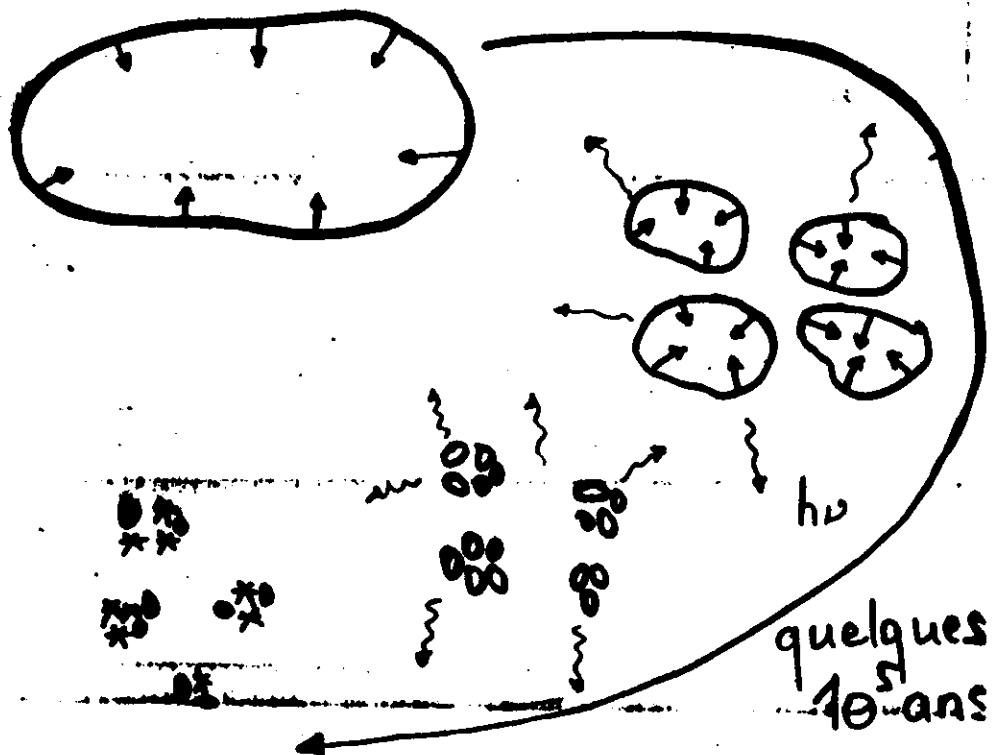
FIG 3. Absorption spectrum towards Cas A. Continuum emission equivalent to 0.22 K has been subtracted from the baseline. The channels are 1 km s^{-1} wide. Note that the absorption corresponds to the velocities of the Orion and Perseus arms at 0 km s^{-1} and -45 km s^{-1} .

image classique:

la fragmentation
est une conséquence
de la contraction

$$M_{\text{Jeans}} \propto \frac{T^{3/2}}{n^{1/2}}$$

support uniquement
thermique



PICTURE OF THE GMC:

- PRIMORDIAL FRAGMENTATION EXISTS BEFORE CONTRACTION
- ASSEMBLY OF ~ 1000 CLOUDS OF 100 M_{\odot} VIRIALISED IN THE GRAVITATIONAL POTENTIAL OF THE GMC.
- TIME SCALE $\gg 10^6$ YEARS
- DENSE CORES HAVE LESS THAN 10% OF THE TOTAL MASS.
- MOST OF THE MASS IS IN THE CLOUDS OF SIZE 1-3 pc
- SCALES ARE SET BY THE EQUATION OF STATE OF ISM \Rightarrow ONLY $\begin{bmatrix} 1-3 \text{ pc} \\ .1 \text{ pc} \end{bmatrix}$ STABLE IF NO NEIGHBOR: COLLAPSE

HIERARCHICAL STRUCTURE IN MOLECULAR CLOUDS

PERAULT
FALGARONE
PUGET

GMC $\sim 10^5 M_{\odot}$

FRAGMENTS $\sim 2 \times 100 M_{\odot}$

DENSE CORES $\sim 2-3 M_{\odot}$

OBSERVATIONS MADE AT BORDEAUX 110-115 GHz
AMHERST 115 GHz
MWO 230 GHz

(68)

not follow a simple $B \propto n^{3/2}$ law, or it may be that we see only the outer parts of the molecular cloud.

By proper star count and a comparison of CO emission, a canonical value $\frac{n_{CO}}{n_{H_2}}$ can be deduced (close to 10^{-4}). A fair amount of carbon is in the form of CO, while the recent detection of the fine structure lines of carbon (at submm wavelength) show that an approximately equal amount is in the form of neutral carbon. The remaining part is probably locked on dust grain (or polymeric molecules).

(69)

B) The Galactic Structure:

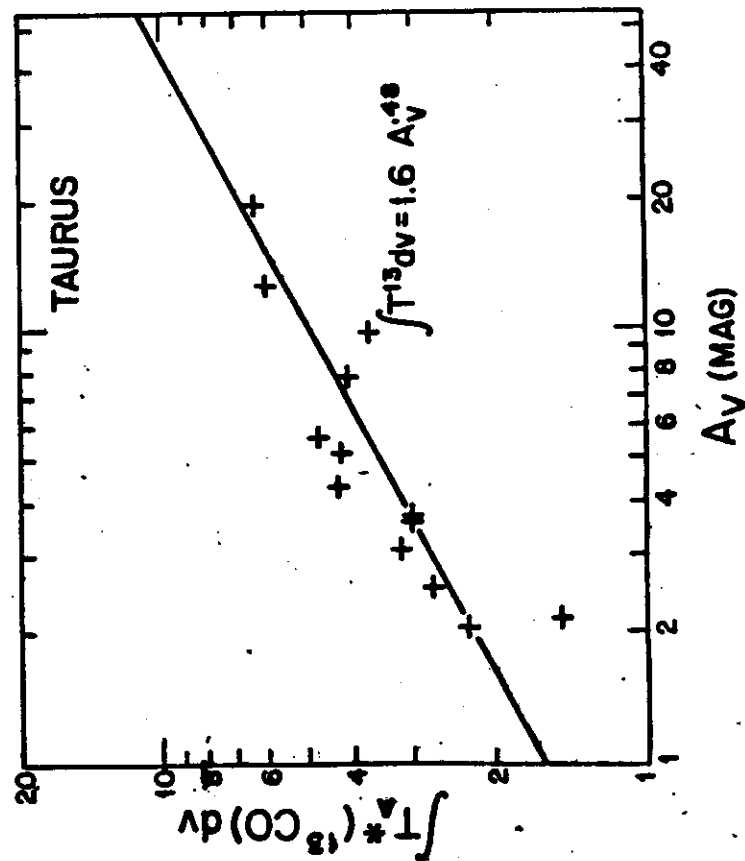
Surveys of the Galactic plane have been made by different groups using 1 to 10 m class telescopes. Of all of them, only part of the sky is fully sampled. The progress in the sensitivity of receivers should permit a complete coverage in the CO 1-0 transition within 2-3 years, while some isotopic survey will be needed later (optical depth effects are not completely understood).

The comparison with HII data show that the galactic structure is well marked in molecular clouds. Except a continuing battle for years (lifetime of the clouds compared the constant of rotation of the Galaxy), it seems now that an agreement has been reached.

The sensitivity of gamma observation is severely limited by the number of photons that the detector sees. It is evident that Giant Molecular Clouds (like the also Ophiuchus complex) are also sources of gamma-ray emission (interaction of cosmic-rays with the dense molecular gas and H₂-decay).

The comparison with IRAS data is complicated by the fact that

part of the emission at $10\ \mu$ is due to polycyclic aromatic molecules, while dust is also associated with HI. As can be seen from the enclosed figure, this picture does not agree well with the previous picture of the dust in our Galaxy (except for the strong infrared sources).



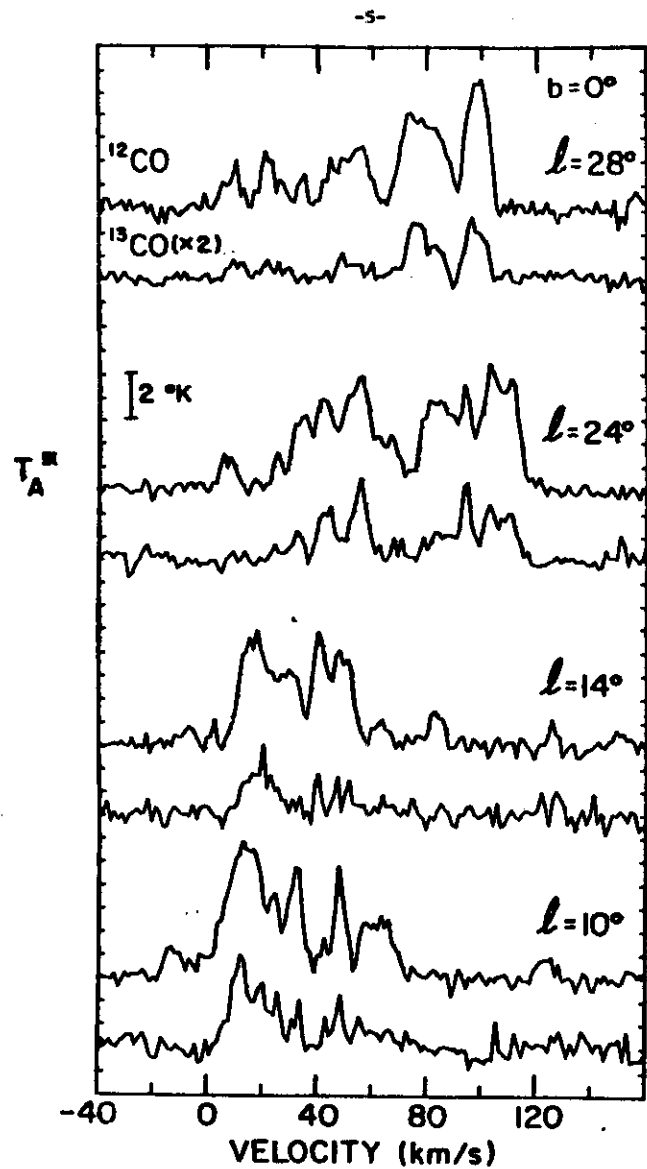


Fig 2. Sample ^{12}CO and ^{13}CO spectra at longitudes 10° , 14° , 24° , 28° . ^{13}CO spectra have been scaled by a factor of 2.

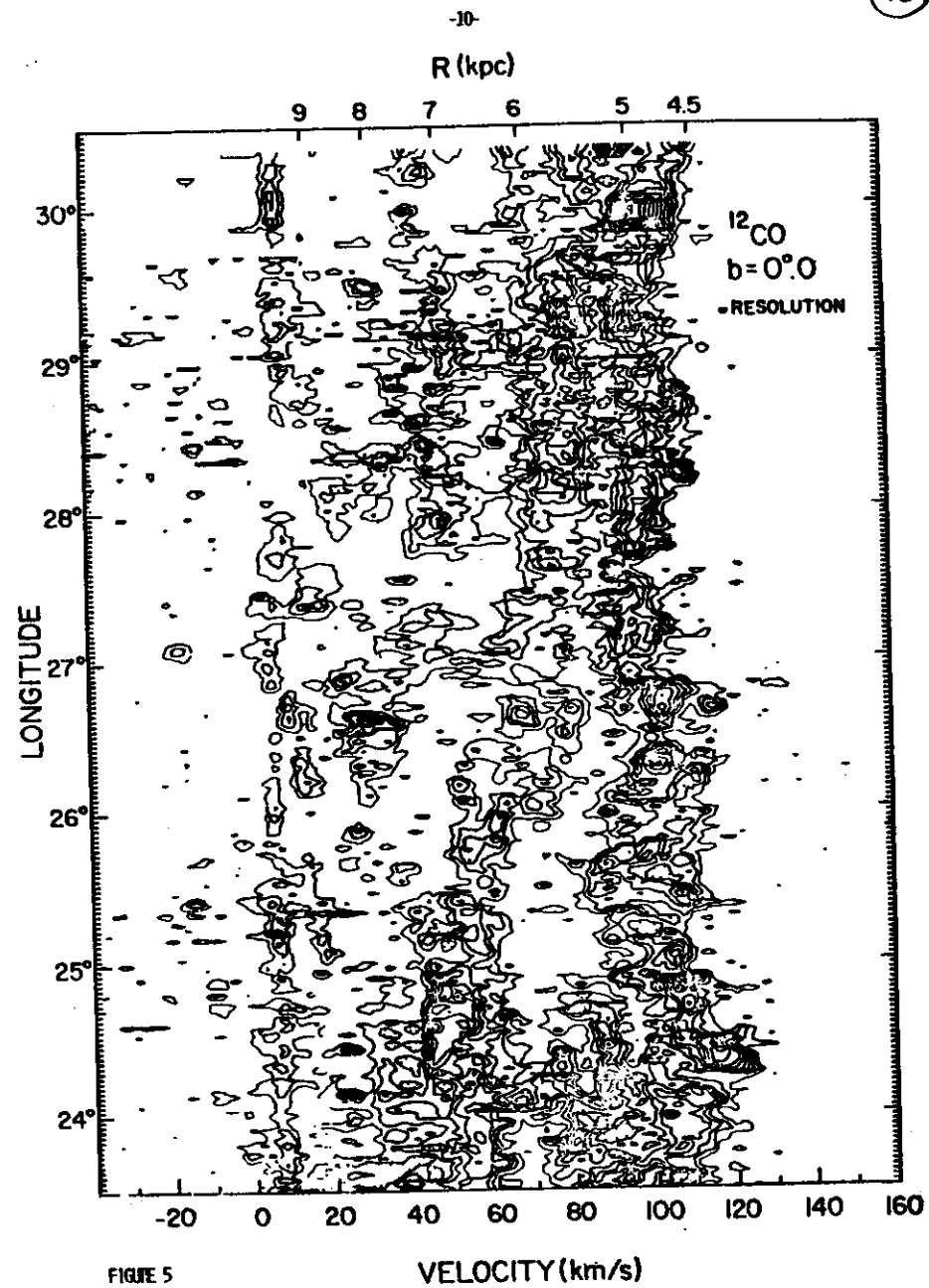
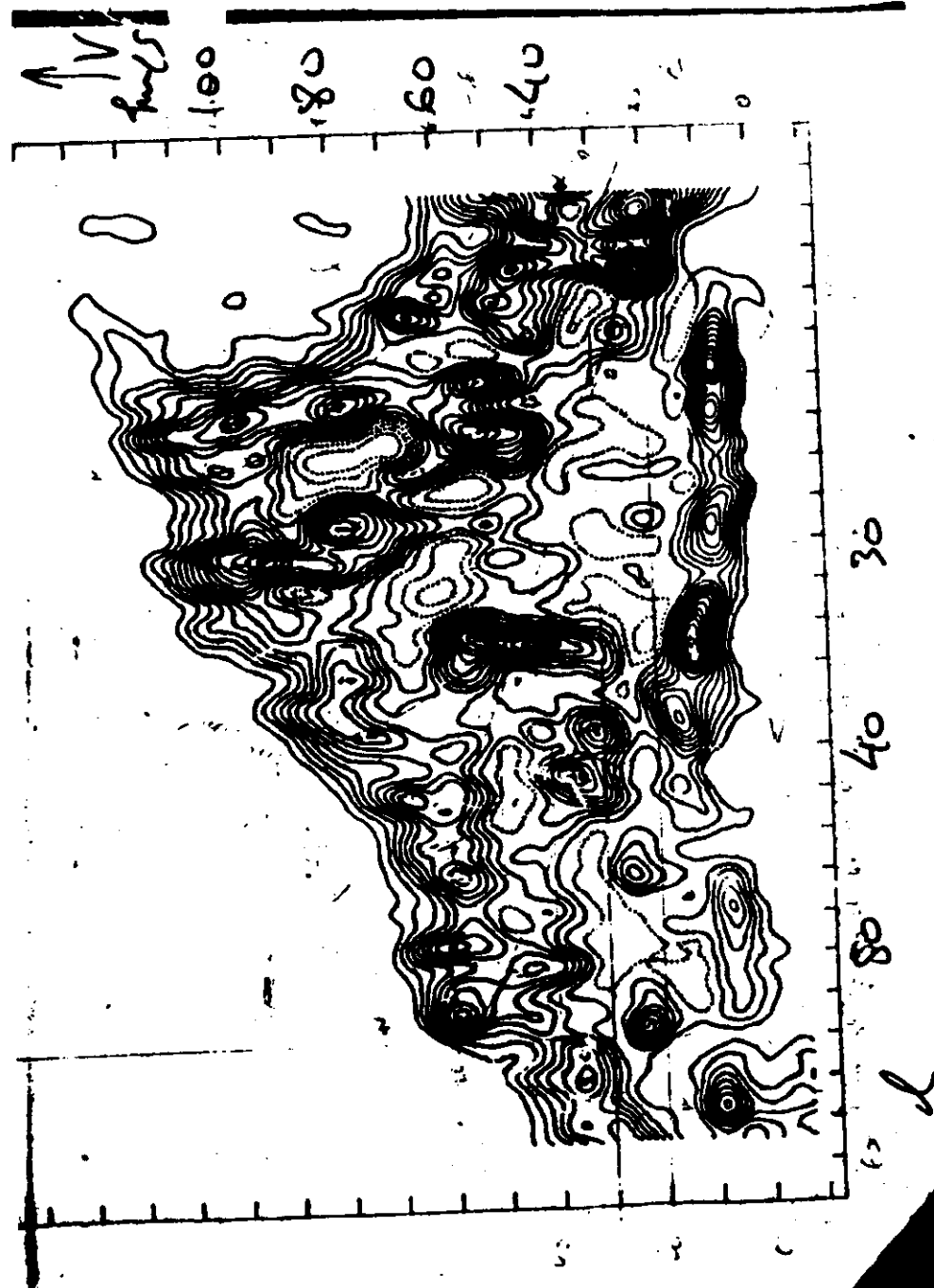
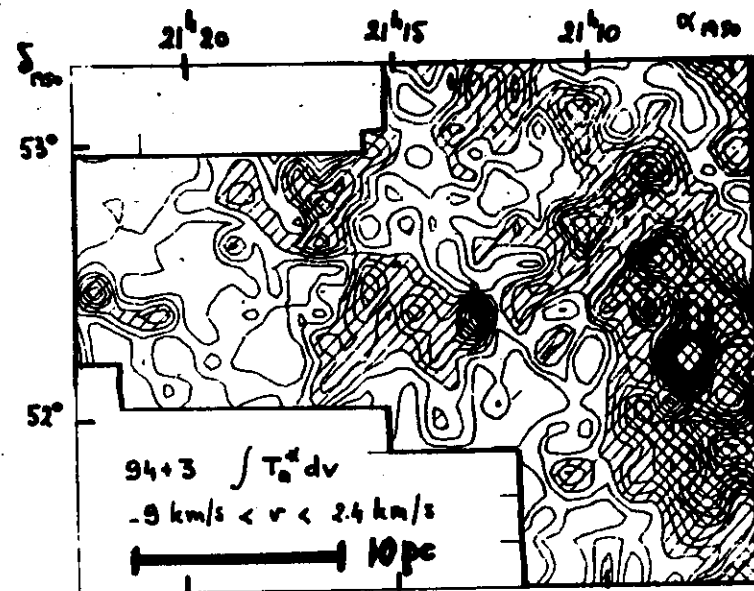


FIGURE 5

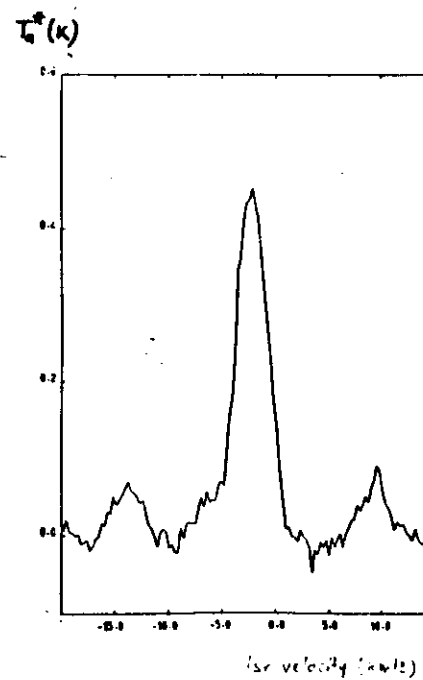


(74)

Fig 1.



(75)



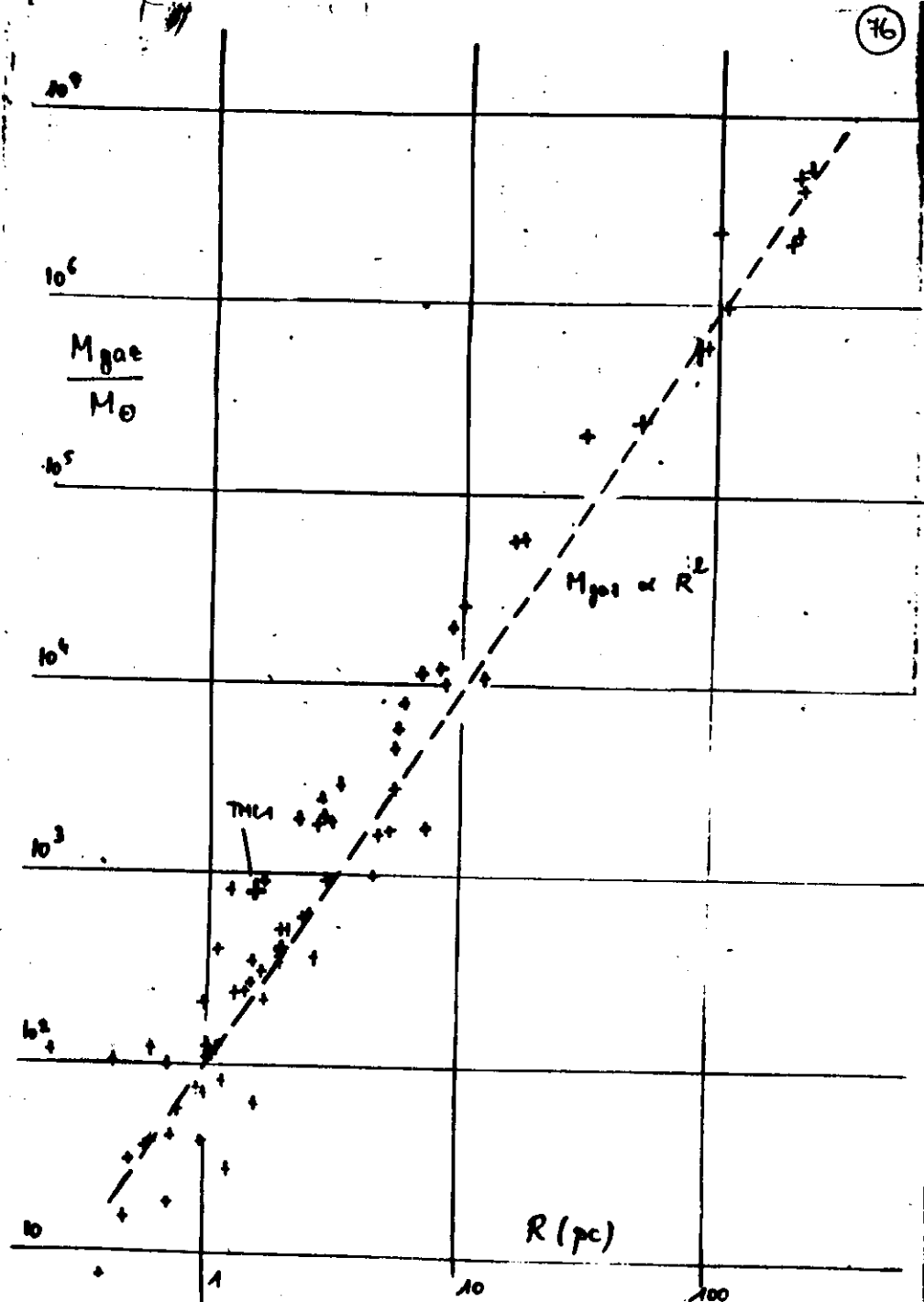
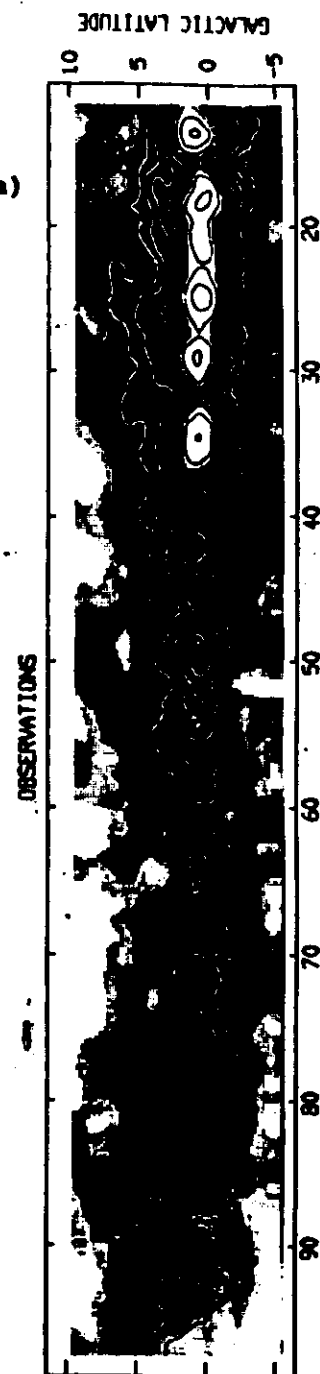
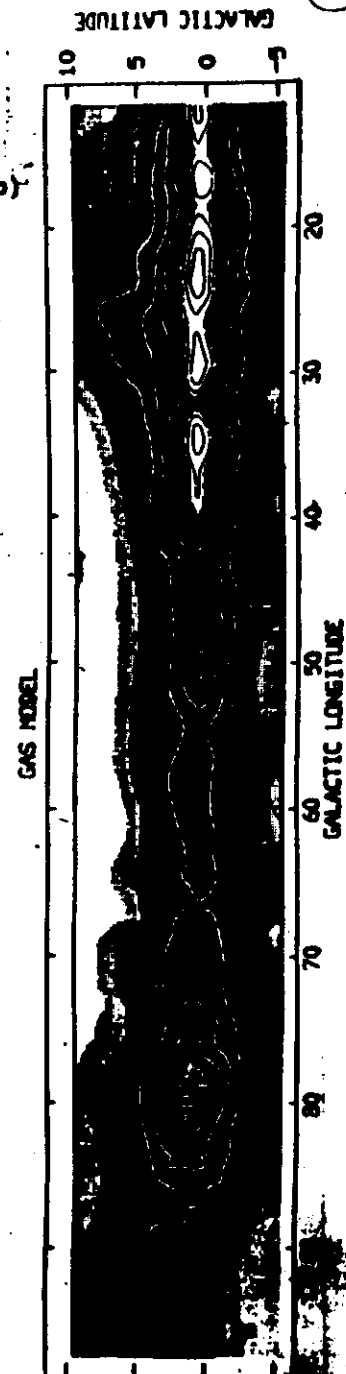


Figure 1

a)



b)



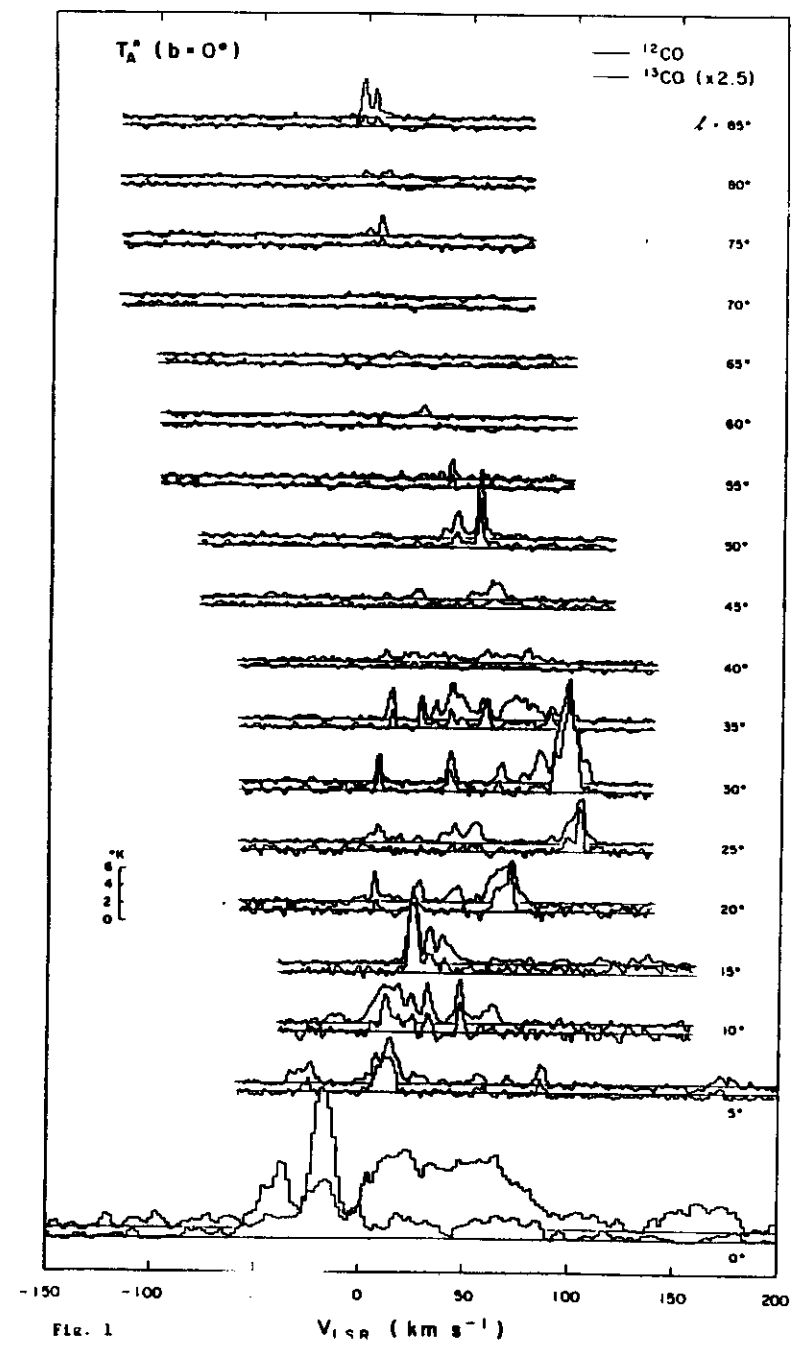


Fig. 1

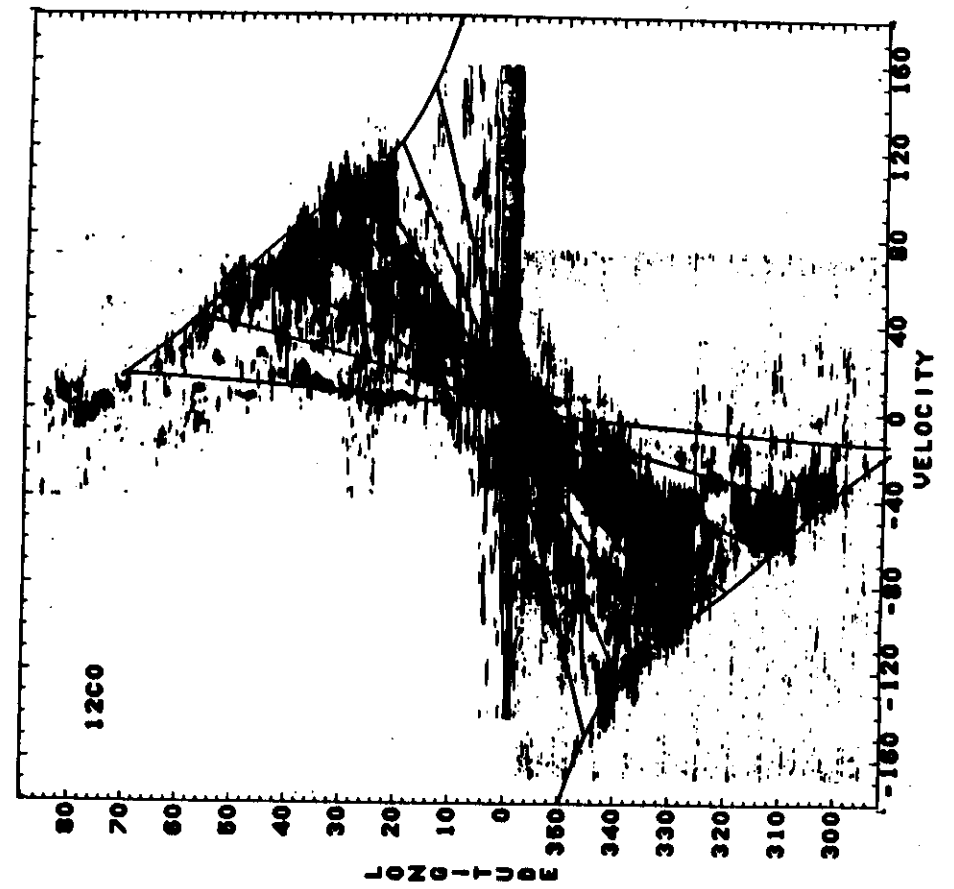
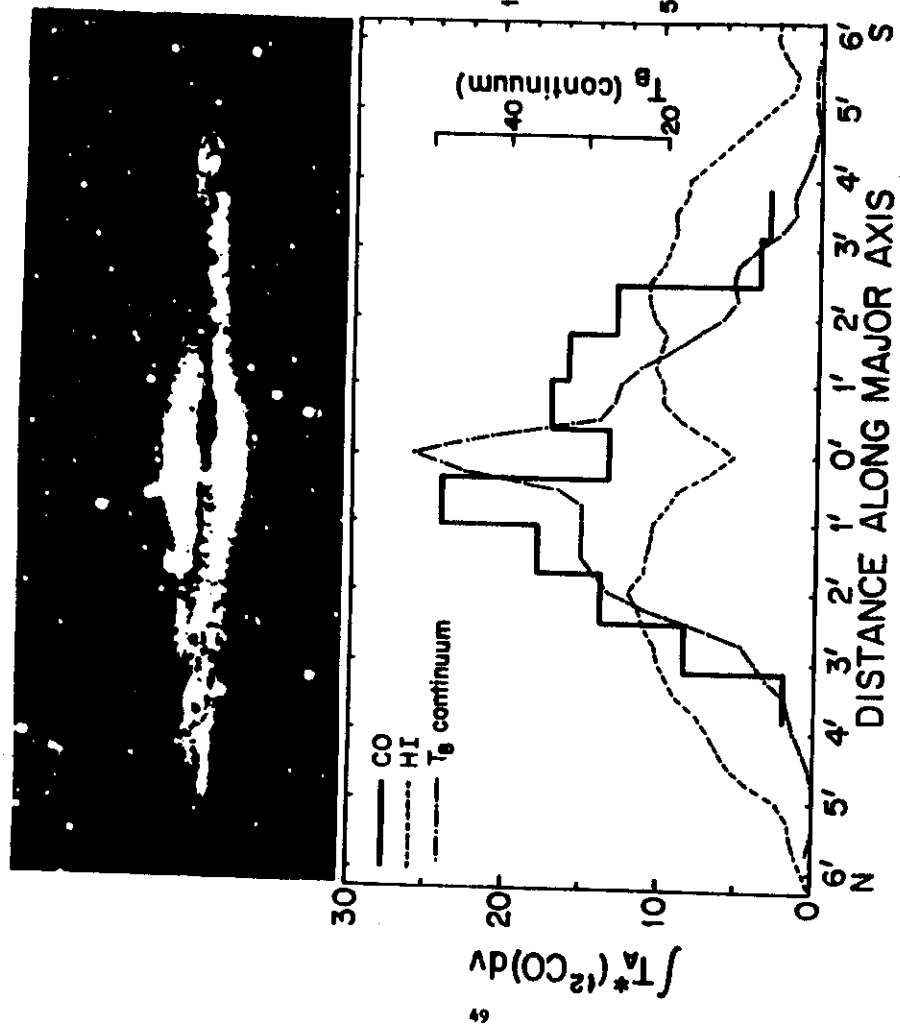
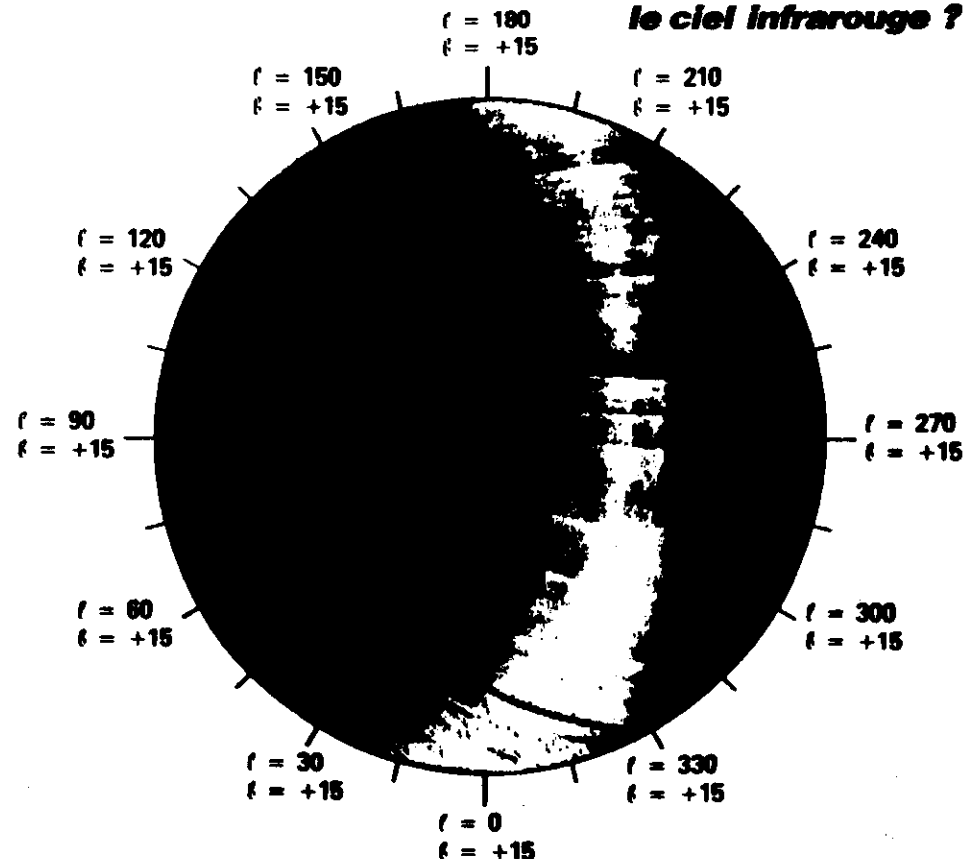


Fig. 2



es poussières microscopiques dans le ciel infrarouge ?



Le 25 janvier 1983 était lancé le premier satellite infrarouge de l'histoire de l'astronomie : IRAS (pour Infrared Astronomical Satellite), fruit d'une collaboration du Royaume-Uni, des Pays-Bas et des États-Unis. Avec son télescope de 57 cm de diamètre, ce satellite permet d'observer le domaine infrarouge dans quatre bandes de longueurs d'onde centrées à 12 μ , 25 μ , 60 μ et 100 μ ⁽¹⁾, domaine où les radiations sont complètement absorbées par l'atmosphère terrestre notamment à cause de la vapeur d'eau (le domaine visible s'arrête à environ 1 μ et celui de la radioastronomie commence à 1 mm).

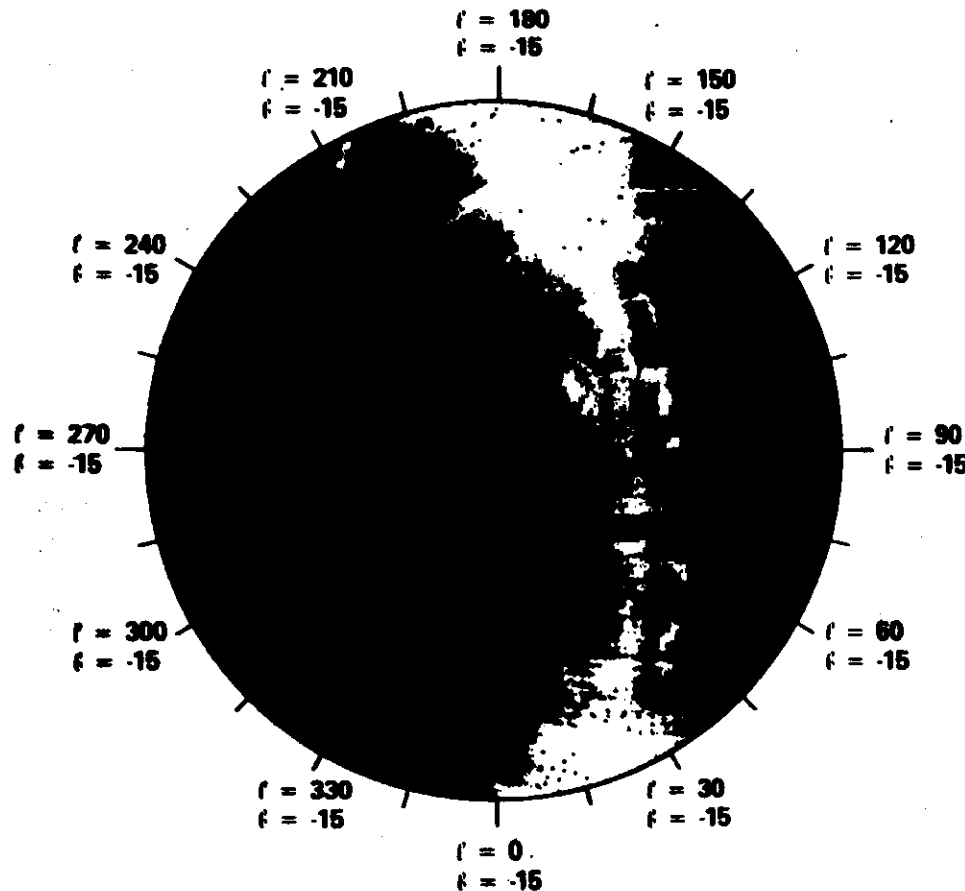
Après dix mois d'observation (il a été arrêté en novembre 1983) IRAS nous a révélé pour la première fois une carte du ciel en infrarouge (voir *la Recherche*, n° 152, p. 254, février 1984). Outre la découverte de systèmes solaires en formation (voir *la Recherche*, n° 150, p. 1596, décembre 1983), IRAS a ouvert la voie à l'étude d'objets et de structures dans l'Univers faits de matière froide et qui n'avaient jamais pu être observés jusqu'alors⁽²⁾. Une de ses découvertes majeures sans doute est la mise en évidence de nouveaux constituants de l'Univers : les cirrus infrarouges, des nuages interstellaires constitués de micro-

particules abondantes en carbone qui tissent une véritable toile d'araignée en tour du système solaire.

On croise de nuages autour du système solaire.

Les nuages de poussières interstellaires peuvent être observés par IRAS car chauffés à de faibles températures, ils émettent principalement de la lumière infrarouge. Tout corps chauffé émet une radiation caractéristique de sa température : plus la température est élevée, plus le maximum du rayonnement émis est déplacé vers les longueurs d'onde-

(1) G. Neugebauer et al., *Ap. J. Letters*, 278, L1, 1984.
(2) Numéro spécial IRAS, *Ap. J. Letters*, 278, mars 1984.



courtes, la relation entre les deux étant donnée approximativement par $T = 2900/\lambda$ ou λ est la longueur d'onde en microns et T la température en degrés Kelvin. Dans la gamme couverte par IRAS (12 à 100 μ) émettent donc essentiellement des corps chauffés à des températures de 30 à 250 K (soit 243 à 23 de nos degrés Celsius).

Or on a pu calculer que, dans la Galaxie, les différents types de rayonnements présents (rayonnement ultraviolet des étoiles, particules de haute énergie du rayonnement cosmique...) étaient suffisants pour chauffer les poussières interstellaires à des températures de l'ordre de

Figure 1. Le satellite d'astronomie infrarouge IRAS a, pendant dix mois en 1983, observé l'ensemble du ciel dans la gamme de longueurs d'onde de 12 à 100 μ ; données inaccessibles aux instruments au sol car entièrement filtrées par l'atmosphère terrestre. A ces longueurs d'onde, les poussières interstellaires chauffées par le rayonnement ultraviolet des étoiles sont la principale source du rayonnement diffus. Sur ces cartes des deux hémisphères célestes Nord (à gauche) et Sud (à droite) est représentée l'intensité de l'émission telle qu'elle a été observée par IRAS (les zones les plus intenses apparaissent en jaune et rouge, les plus faibles en bleu et noir). Chaque carte est couverte par chacun des piliers Nord et Sud de la Galaxie et produite en longitude (l) et latitude (b) galactiques. La bande sombre et étroite (de $l = 90^\circ$ à $l = 270^\circ$) correspond à des régions qui n'ont pu être observées. A l'exception des nébuleuses d'Orion et de la Carina qui apparaissent comme deux taches brillantes (vers $l = 210^\circ$ et $l = 265^\circ$), le reste de l'émission est remarquablement uniforme. A la grande surprise des chercheurs, les nuages de poussières qui apparaissent ainsi ou noir se correspondent pas aux concentrations de matière interstellaires connues jusqu'à. Ils se répartissent uniformément, couvrant le ciel infrarouge de structures filamenteuses : des cirrus infrarouges d'aspect analogue aux cirrus de l'atmosphère terrestre. Ces structures contingentes de l'Univers sont encore très énigmatiques. Les caractéristiques de leur émission infrarouge suggèrent une composition de micrograins à forte teneur en carbone qui pourraient être en fait un signal de grosses molécules cycliques. Leur origine reste encore mystérieuse et leur distance exacte est encore inconnue. (Cliché JPL).



Figure 2. Les sources intenses de rayonnement infrarouge sont les poussières interstellaires chauffées par le rayonnement ultraviolet des étoiles. Dans la Galaxie, les nuages moléculaires (des nuages où la densité du gaz d'hydrogène interstellaire est suffisante pour recombiner l'hydrogène atomique en molécules H_2) contiennent une forte proportion de poussières. Sur cette photo prise par IRAS à la longueur d'onde de 12 μ , dans une région du ciel de 25° par 25° , le nuage moléculaire R Carinae Australis apparaît clairement. L'intensité du rayonnement infrarouge émis est représentée par une échelle de couleurs (indiquée au bas de la figure). Le rayonnement infrarouge observé par IRAS pour plusieurs de ces nuages est trop étalé pour être émis par des poussières de taille « normale » (0,1 à 1 micron) chauffées par le rayonnement galactique ambiant. Il pourrait, en revanche, s'expliquer si une certaine proportion de la poussière dans ces nuages moléculaires était de nature type que celle suggérée par les cirrus infrarouges, c'est-à-dire des micro-grains de quelques 0,005 micron de diamètre. (Cliché A. Lema et P. Vassiliou).

25 à 30 K. Un unique type de poussières était connu jusque-là : des grains de silicate d'une taille de 0,01 à 1 micron mélangés au gaz de la Galaxie et distribués approximativement comme lui, c'est-à-dire concentrés dans la plus galactique. La cartographie du ciel établie par IRAS à la longueur d'onde de 100 μ aurait donc dû révéler ces poussières approximativement selon le dessin de la Galaxie.

Or, à la grande surprise des chercheurs, l'émission observée à cette longueur d'onde n'est pas particulièrement localisée, mais semble venir de toutes les directions (fig. 1). Elle couvre l'ensemble du ciel infrarouge de structures filamenteuses à la manière de certains nuages terrestres qui sont les cirrus, ce qui leur a

valé d'être baptisés « cirrus infrarouges ».

A cause de l'uniformité de leur répartition, ces cirrus correspondent sans doute à de la matière interstellaire proche qui, comme un cocon, entoure le système solaire. Il est impossible à l'heure actuelle de leur plus précisément leur distance. Il a été suggéré qu'une partie de ces nuages soit en fait très proche et située aux confins du système solaire (à une distance inférieure à une année-lumière), mais la majorité est sans doute interstellaire à des distances approximatives de 600 années-lumière.

Dans l'étude de ces cirrus infrarouges, une deuxième surprise allait venir des cartes établies par IRAS aux trois autres longueurs d'onde. Ces nuages apparais-

sent en effet de façon très claire sur la carte à 60 μ , et F. Boulanger de l'Observatoire de Meudon, actuellement à l'université de Groningen, a découvert qu'ils émettaient également un rayonnement à 12 et 25 μ ! L'émission à ces longueurs d'onde, bien que quatre fois plus faible que celle à 160 μ , est beaucoup plus importante que celle que l'on pouvait envisager pour la poussière interstellaire.

De la poussière chaude
dans des étoiles.

Cette émission était d'abord pensée insoupçonnée à cause de la lumière solaire, des aux poussières qui existent également à l'intérieur du notre système solaire. Du fait de l'attraction du Soleil, ces pou-

C/ Extragalactic Molecular Astronomy: a field in its infancy

Since its discovery in 1975, 200 galaxies have been observed and approximately 60 detected. The main problems associated with these observations are the lack of sensitivity

lack of resolution

The main beam is:

BTL	1.7'
KPNO	1'
UMASS	50"
Swedish	33"
IRAM 30m	22"
Nobeyama 45m	15"
Cal Tech 30m	8"

Selection effects are strongly affecting the data:

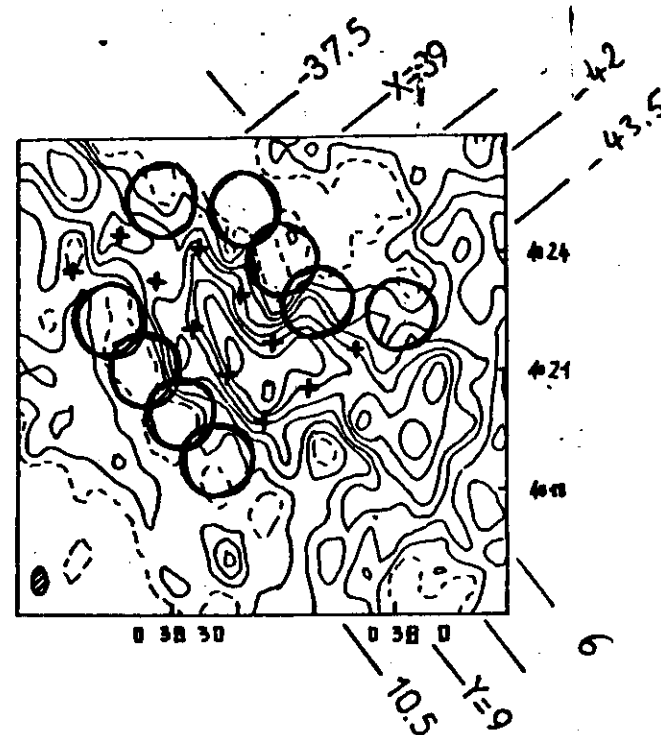
- inclination of the galaxies
- dilution in velocity

The advent of more sensitive receivers and larger collecting areas should suppress this bias.

The main results can be summarized as follows:

- radial distribution of CO/HI
 - $n(\text{H}_2)/n(\text{HI})$
 - Spiral Structure
 - Size of the Giant Molecular Clouds
- } only in 1731

Isotopically substituted molecules have been observed for the first time in 1731 (^{13}CO) and in 1782 (C^{18}O). The ratios to the main line is difficult to analyze.



1731

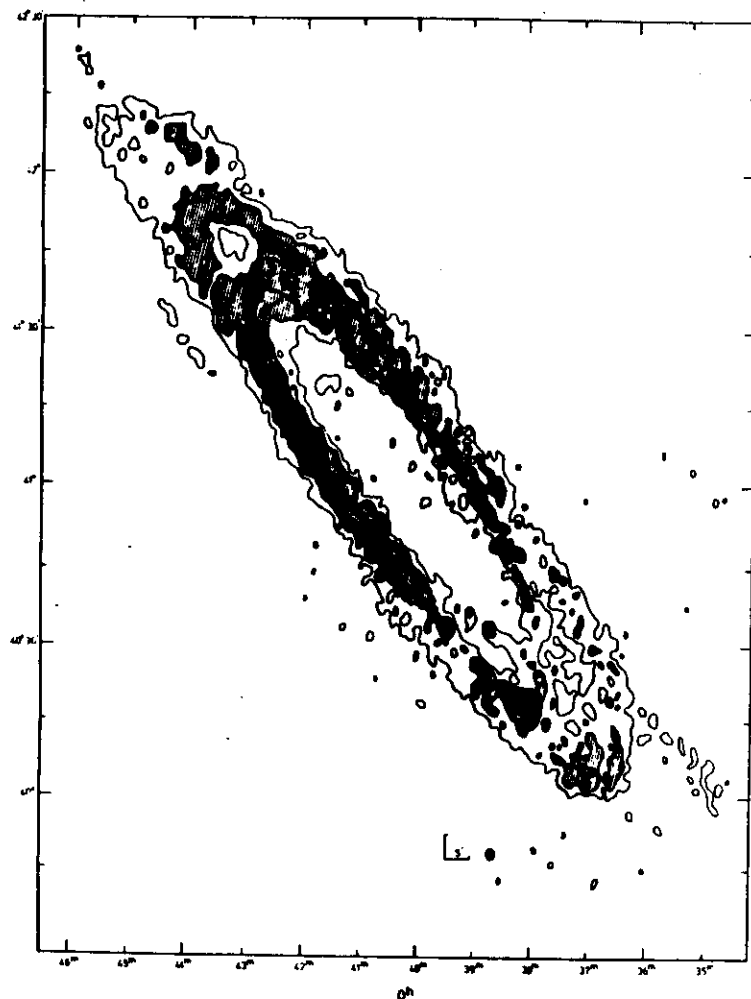
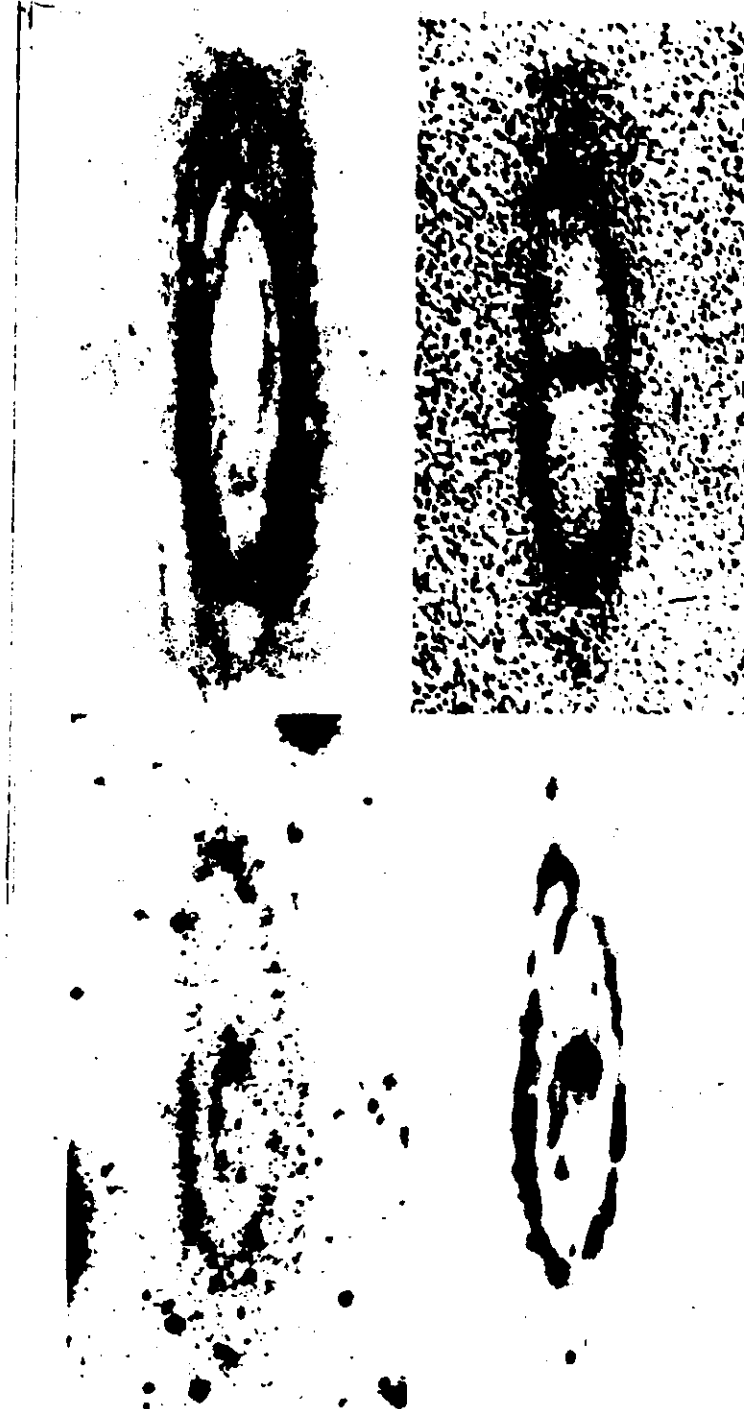
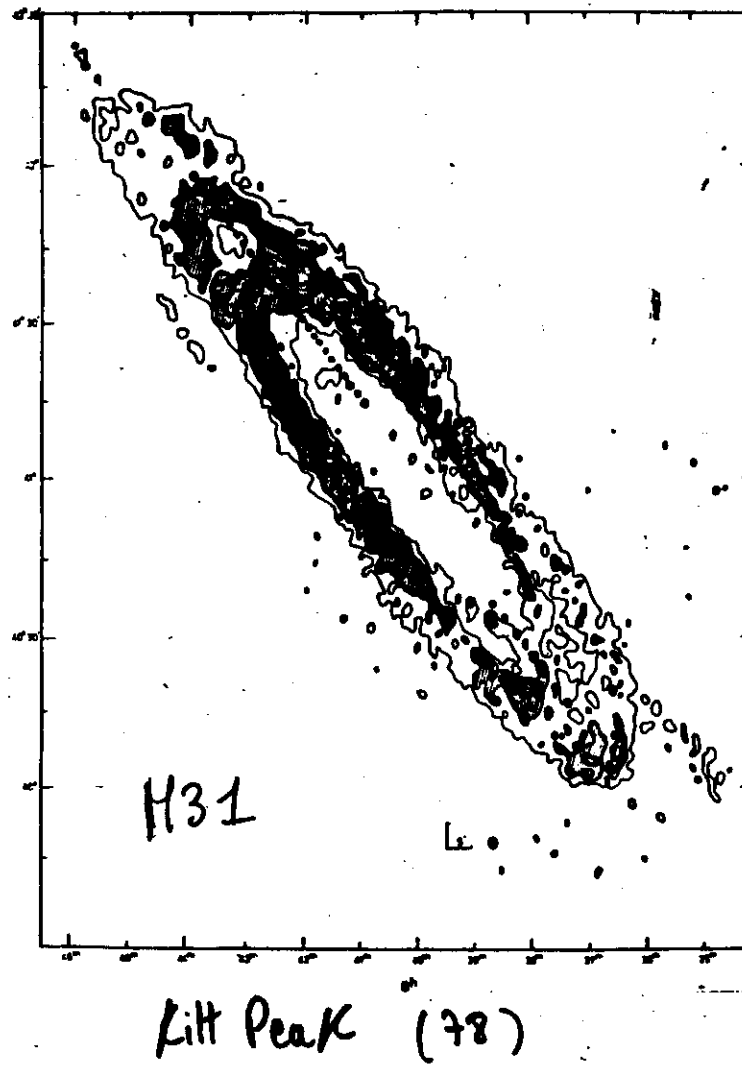
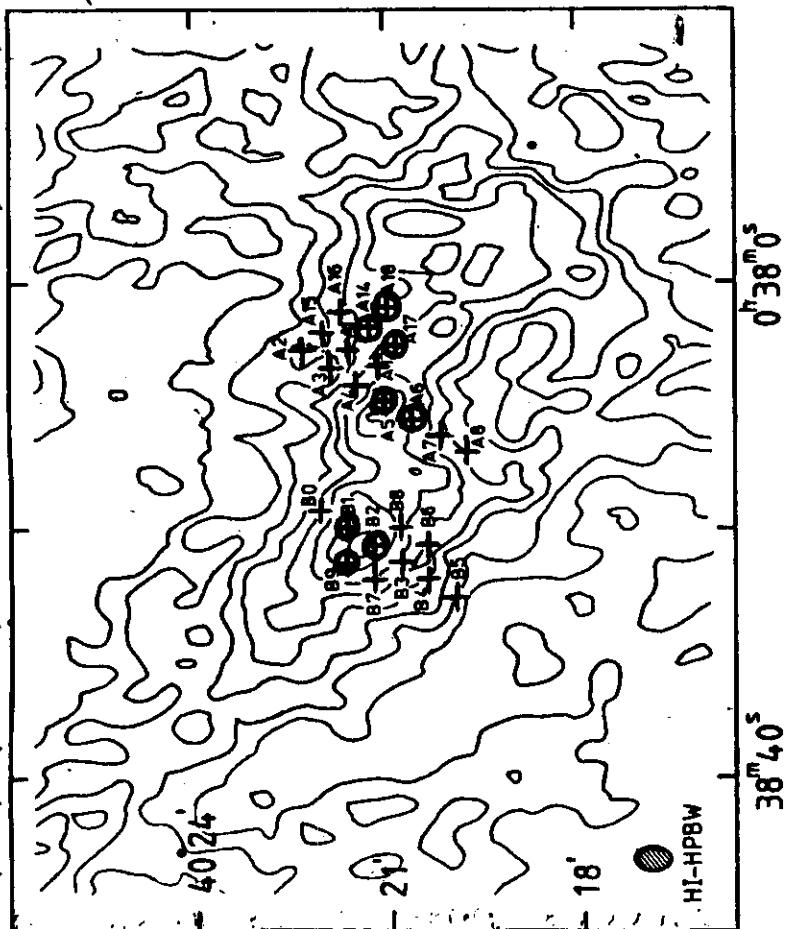


FIG. 1. Integrated HI emission from M31. The beam size of 1.5 x 2.2 arc min is shown as a shaded ellipse. The contour interval is 600 K km s⁻¹, with regions above the 1200 K km s⁻¹ level, and also the 2400 K km s⁻¹ level, being shown shaded. The so-called "SW companion" of M31 is not shown, being beyond the SW extension of this map.

86



87



(90)

Galaxies cartographiées en CO

	Type	Anneau	i	Nbre de lobes	D Mpc
M31	Sb	⊗	77	24000	.7
N253	Sc AB		79	160	2.5
M33	Scd	points	55	3100	.7
N891	Sb	?	88	9	9.5
N1068	Sb		43	50	16
N2403	Scd AB	points	60	230	3.2
N2841	Sb	⊗	68	36	9
N2903	Sbc AB	3 points	70	45	6
M82	Irr	filament.	66	43	3.2
N3627	Sb AB		60	55	9
N3628	Sb pec		76	77	9
N4254	Sc		27	9	20
N4258	Sbc AB		72	50	5
N4321	Sbc AB		35	55	20
M51	Sbc		20	71	7.6
M83	Sc AB		24	100	4.3
N5248	Sbc AB	⊗	55	20	15
M101	Scd AB		22	590	8.5
N6946	Scd AB		30	90	4.6
N7331	Sbc	⊗	75	43	15
IC342	Scd AB		25	415	3
NGC 2	Sb		60	43	4

(91)

NGC 5236 (M83)

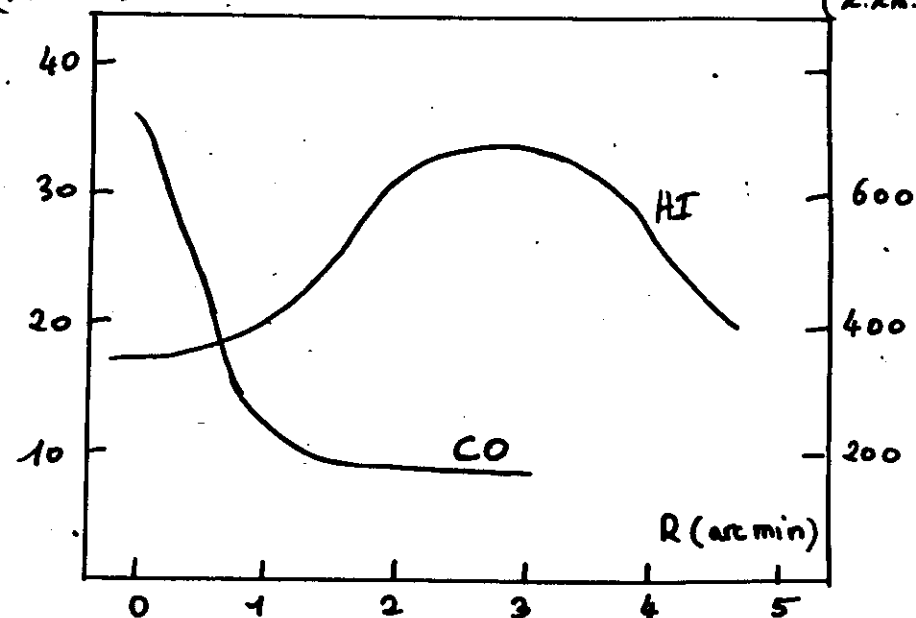
from Combes et al (1978)

$$\int T_A(\text{CO}) dV$$

(K.Km.s⁻¹)

$$\int T_B(\text{HI}) dV$$

(K.Km.s⁻¹)



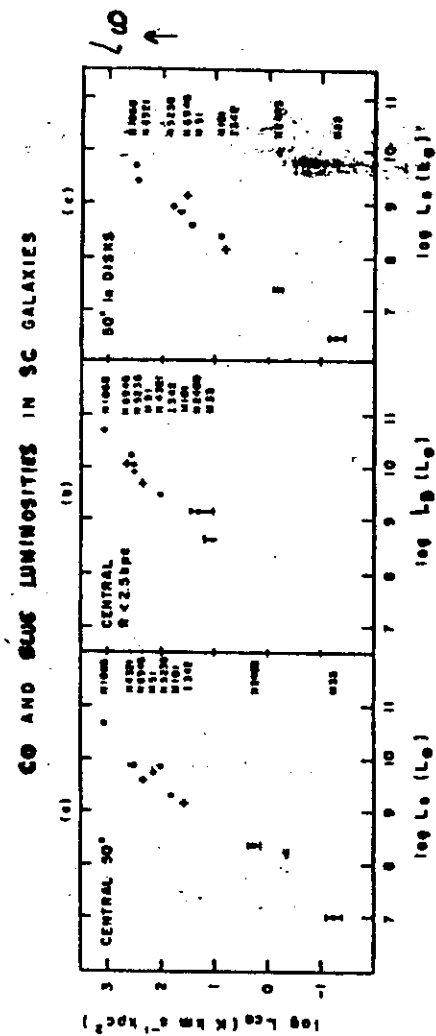
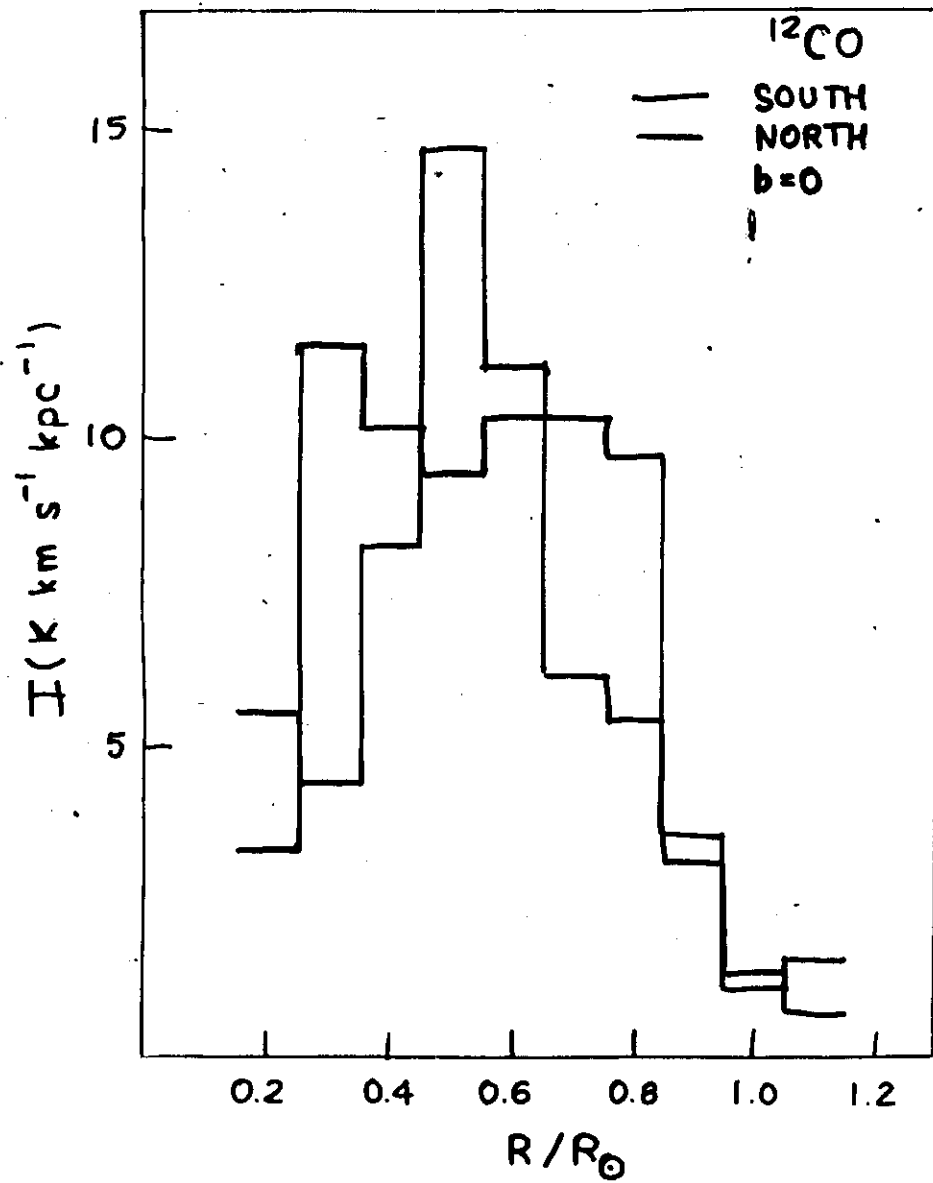


Figure 7. Comparison of CO and blue luminosities in (a) the central 50", (b) regions 50" in diameter centered on the nuclei, and (c) regions 50" in diameter in the disks for nine galaxies ($L_{\text{CO}} \pm 10\%$ Area Beam). Over four orders of magnitude in luminosity, a linear correlation is evident. The open circle represents the Seyfert galaxy NGC 1068 which has the highest luminosity. The uncertainties in L_{CO} are given for M33 and NGC 2043, and for the other galaxies are smaller than the dots plotted. Distance uncertainties do not alter the correlation since both luminosities depend on D^2 . The best fit to the data in (b) is $L_B = 4 \times 10^7 L_{\text{CO}}^{1.0}$.

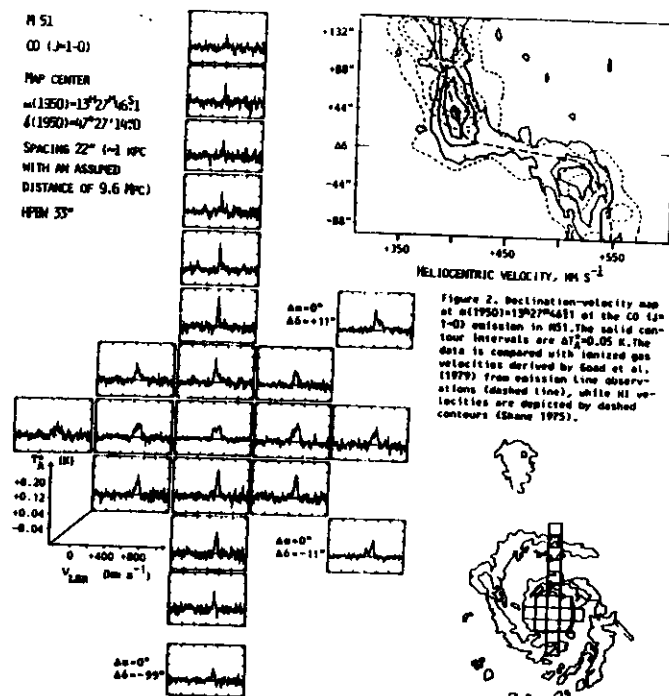


Figure 1. CO emission in M51. The inset diagram shows the position of the spectra on a tracing of the optical outline of the galaxy. The intensity scale is given in antenna temperature corrected for radome and atmospheric attenuation. The main beam efficiency is 0.3 and the mean efficiency is 0.4. To get from V_{LSR} to heliocentric velocity, subtract 11.7 km s^{-1} .

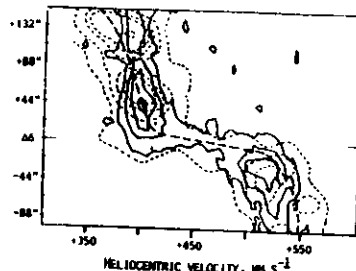


Figure 2. Declination-velocity map of M51. The solid contours are $\Delta T_{\text{MB}} = 0.05 \text{ K}$. The data is compared with ionized gas velocities derived by Sand et al. (1979) from emission line observations (dashed line), while HI velocities are depicted by dashed contours (Sand 1975).

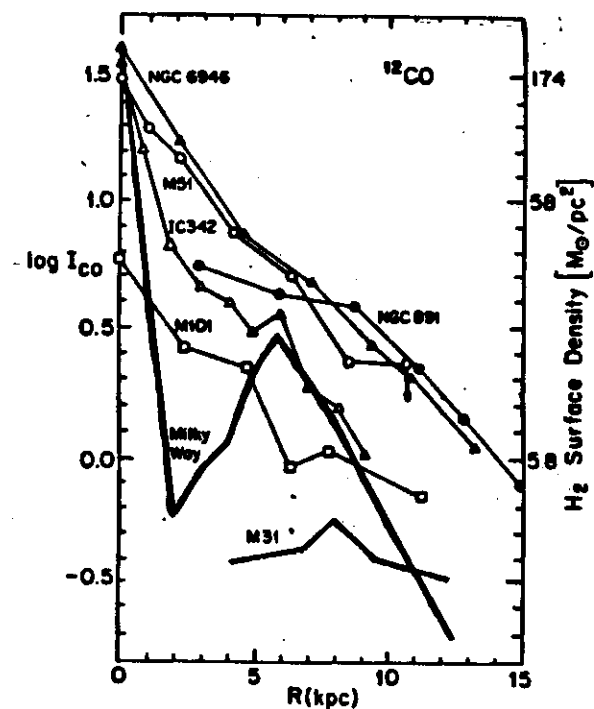


FIG. 9. RADIAL DISTRIBUTION OF CO EMISSION IN 6 GALAXIES COMPARED WITH THE MILKY WAY.

MILLIMETER RADIO TELESCOPES:

		$\phi(m)$	λ_{mm}
<u>SINGLE DISH:</u>	EFFELSBERG	100	8
	NOBEYAMA	45	3
	GRENADA	30	1.5
	GOTEBORG	21	3
	ATIBAIA	14	5
	FINLAND	14	3
	SPAIN	14	5
	KPNO	12	1.3
	AMHERST	14	3
	CAL TECH	10	1
	BANGALORE	10	1
	HAWAII	10	3

INTERFEROMETERS:

IRAM	3x15m	.7
NOBEYAMA	5x10	1
CAL TECH	3x10	3
BERKELEY	3x6	3

MAJOR ROLE OF SMALL INSTRUMENTS:

COLUMBIA	1.2m	3
BORDEAUX	2.5m	3
TOKYO	1m	3
BERKELEY	6m	3
MWO	5m	1

