

Sternentstehung - Star Formation

Sommersemester 2009

Henrik Beuther

3.4 Today: Introduction & Overview

10.4 Karfreitag

17.4 Physical processes I

24.4 Physical processes II

1.5 Tag der Arbeit

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22.5 Accretion disks

29.5 No lecture

5.6 Jet theory (Christian Fendt)

12.6 Observations of outflows and jets

19.6 Protostellar evolution, stellar birthline, pre-main sequence evolution

26.6 High-mass star formation, clusters and the IMF

3.7 Extragalactic star formation (Eva Schinnerer)

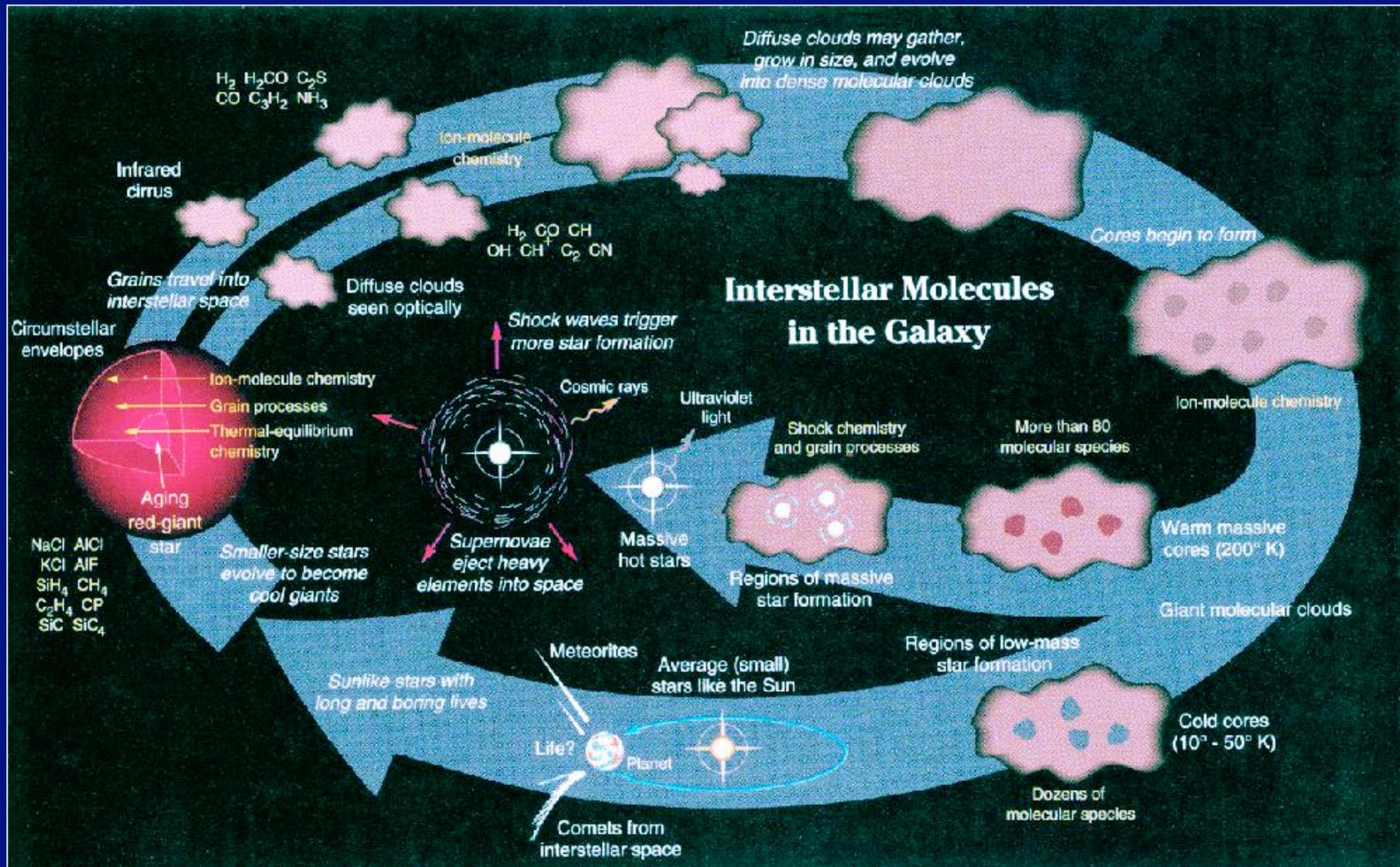
10.7 Summary

More Information and the current lecture files: http://www.mpia.de/homes/beuther/lecture_ss09.html
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Topics today

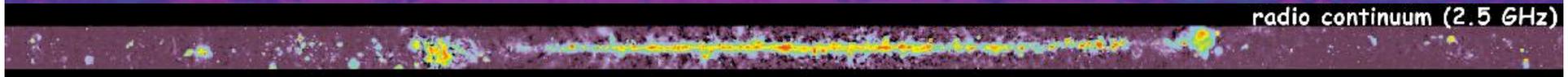
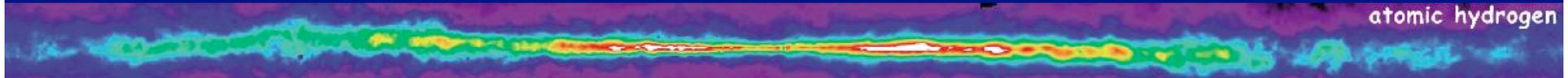
- Molecules and depletion
- Heating and cooling
- Radiation transfer and column density determination
- Line profiles
- Applications

The cosmic cycle

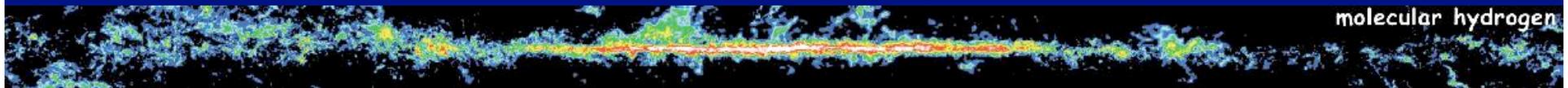


Basics I

Neutral and ionized medium



Stars form in the dense molecular gas and dust cores



Most important astrophysical tools:

Spectral lines emitted by various molecules

Absorption and thermal emission from dust

Basics II

High dust column densities block optical and UV-light in dark cores

--> important requirement that molecules can form and survive

History:

- Late 1930s: Detection of CH, CH⁺ and CN in diffuse clouds by absorption of optical light by background stars

- 1960s: Detection of OH, NH₃ and H₂O at radio wavelength

Formation of molecules is an energy problem, two atoms approach each other with positive total energy, hence would simply rebound if no energy could be given away somehow ... Possibilities:

- Simultaneous collision with 3rd atom carrying away energy

--> unlikely at the given low densities

- Form a molecule in excited state, and then radiating away energy

--> probability of such radiative association low as well

- Neutral-neutral reactions from unstable "activated complexes" (intermediate between reactants and products) which may form larger molecules

--> activation barrier ~100K, impossible on cold dark cores

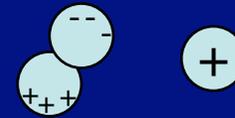
- **Ion-molecule or ion-atom reactions can solve energy problem**

- **Neutral-neutral reactions on dust grain surfaces (catalytic) important**

Basics III

Ion induces dipole moment in atom or molecule which creates an electrostatic attraction between the two.

--> effective cross section increases over geometric values



Even at low temperatures such reactions can account for large fraction of interstellar molecules. However, there are not enough ions available at any time to account for large H₂ abundances

--> grain surface chemistry important

Simple molecules like CO or CS explainable by ion-molecule chemistry, less clear for complex molecules, grain surface chemistry important

No larger inorganic species than, e.g., NH₃, however large organic molecules --> carbon bond important in interstellar chemistry

CO most abundant molecule after H₂

--> most information about molecular clouds from CO observations

Molecules in Space

2	3	4	5	6	7	8	9	10	11	12	13 atoms
H ₂	C ₃	c-C ₃ H	C ₅	C ₅ H	C ₆ H	CH ₃ C ₃ N	CH ₃ C ₄ H	CH ₃ C ₅ N?	HC ₉ N	CH ₃ OC ₂ H ₅	HC ₁₁ N
AlF	C ₂ H	l-C ₃ H	C ₄ H	l-H ₂ C ₄	CH ₂ CHCN	HCOOCH ₃	CH ₃ CH ₂ CN	(CH ₃) ₂ CO			
AlCl	C ₂ O	C ₃ N	C ₄ Si	C ₂ H ₄	CH ₃ C ₂ H	CH ₃ COOH?	(CH ₃) ₂ O	NH ₂ CH ₂ COOH?			
C ₂	C ₂ S	C ₃ O	l-C ₃ H ₂	CH ₃ CN	HC ₅ N	C ₇ H	CH ₃ CH ₂ OH	CH ₃ CH ₂ CHO			
CH	CH ₂	C ₃ S	c-C ₃ H ₂	CH ₃ NC	HCOCH ₃	H ₂ C ₆	HC ₇ N				
CH ⁺	HCN	C ₂ H ₂	CH ₂ CN	CH ₃ OH	NH ₂ CH ₃	CH ₂ OHCHO	C ₈ H				
CN	HCO	CH ₂ D ⁺ ?	CH ₄	CH ₃ SH	c-C ₂ H ₄ O	CH ₂ CHCHO					
CO	HCO ⁺	HCCN	HC ₃ N	HC ₃ NH ⁺	CH ₂ CHOH						
CO ⁺	HCS ⁺	HCNH ⁺	HC ₂ NC	HC ₂ CHO							
CP	HOC ⁺	HNCO	HCOOH	NH ₂ CHO							
C ₅ i	H ₂ O	HNCS	H ₂ CHN	C ₅ N							
HCl	H ₂ S	HOCO ⁺	H ₂ C ₂ O	HC ₄ N							
KCl	HNC	H ₂ CO	H ₂ NCN								
NH	HNO	H ₂ CN	HNC ₃								
NO	MgCN	H ₂ CS	SiH ₄								
NS	MgNC	H ₃ O ⁺	H ₂ COH ⁺								
NaCl	N ₂ H ⁺	NH ₃									
OH	N ₂ O	SiC ₃									
PN	NaCN	C ₄									
SO	OCS										
SO ⁺	SO ₂										
SiN	c-SiC ₂										
SiO	CO ₂										
SiS	NH ₂										
CS	H ₃ ⁺										
HF	SiCN										
SH	AlNC										
FeO(?)	SiNC										

About 150 detected interstellar molecules as of April 2009 (www.cdms.de).
36 (+2 tentative) molecular detection in extragalactic systems.

A few important molecules

Mol.	Trans.	Abund.	Crit. Dens. [cm ⁻³]	Comments
H ₂	1-0 S(1)	1	8x10 ⁷	Shock tracer
CO	J=1-0	8x10 ⁻⁵	3x10 ³	Low-density probe
OH	² Π _{3/2} ; J=3/2	3x10 ⁻⁷	1x10 ⁰	Magnetic field probe (Zeeman)
NH ₃	J,K=1,1	2x10 ⁻⁸	2x10 ⁴	Temperature probe
CS	J=2-1	1x10 ⁻⁸	4x10 ⁵	High-density probe
SiO	J=2-1		6x10 ⁵	Outflow shock tracer
H ₂ O	6 ₁₆ -5 ₂₃		1x10 ³	Maser
H ₂ O	1 ₁₀ -1 ₁₁	<7x10 ⁻⁸	2x10 ⁷	Warm gas probe
CH ₃ OH	7-6	1x10 ⁻⁷	1x10 ⁵	Dense gas/temperature probe
CH ₃ CN	19-18	2x10 ⁻⁸	2x10 ⁷	Temperature probe in Hot Cores

Basics IV

Depletion of molecules on dust grains

The grains are moving at v_{therm} relative to molecules

$$E = 1/2 m v_{\text{therm}}^2 = 3/2 k_b T \Rightarrow v_{\text{therm}} = (3k_b T/m)^{1/2}$$

n grains sweeps out cylindrical volume in time Δt of
 $n(\pi a^2)v_{\text{therm}}\Delta t$ (a: grain radius)

Probability of molecule in volume V to be struck by grain in time Δt

$$P(\Delta t)/V = n(\pi a^2)v_{\text{therm}}\Delta t / V$$

Hence the collision time t_{coll}

$$t_{\text{coll}} = 1/(n(\pi a^2)v_{\text{therm}}) = 1/(n_H \Sigma v_{\text{therm}}) \quad (n_H: \text{density}; \Sigma: \text{cross section})$$

For example CS: $v_{\text{therm}} \sim 5 \times 10^3 \text{ cm s}^{-1}$ at 10K, and $n_H \sim 10^4 \text{ cm}^{-3}$
 $t_{\text{coll}} \sim 6 \times 10^5 \text{ yr}$

Depletion time-scale very short

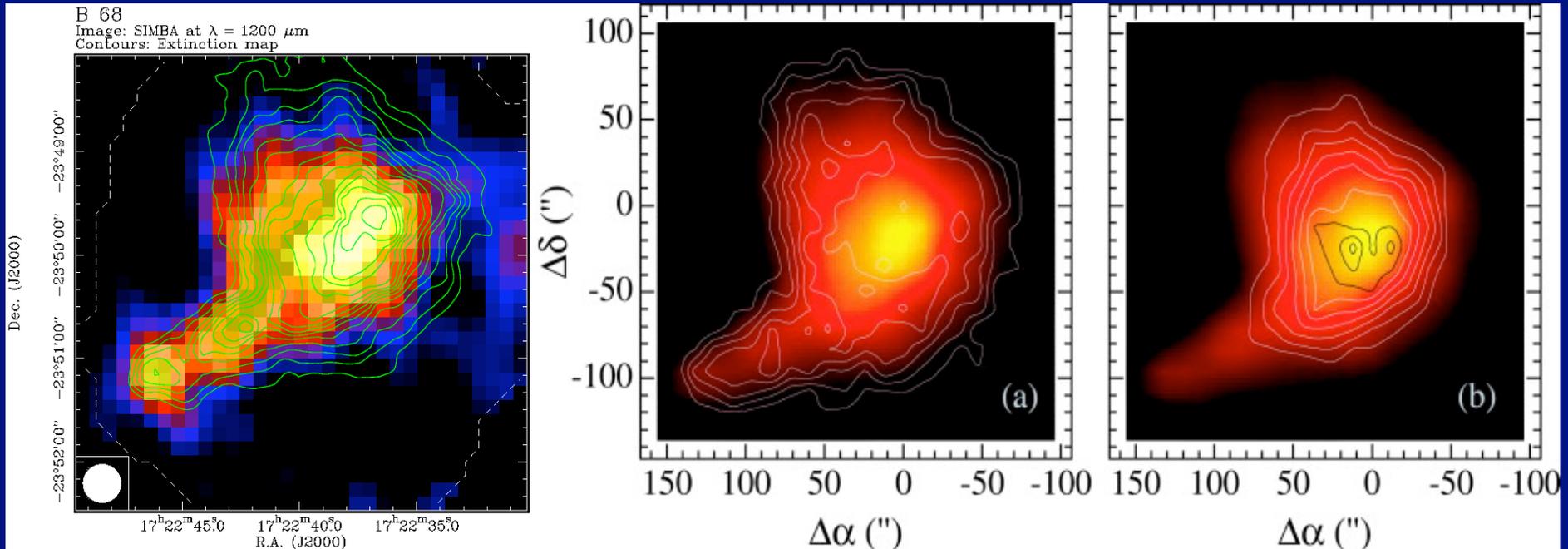
--> mechanisms for reinjecting molecules from grains important

Depletion example

1.2 mm Dust Continuum

$C^{18}O$

N_2H^+



Possible mechanisms working against depletion:

- UV radiation (not working in dense cores)
- In small grain, heat from chemical grain surface reactions could raise temperature
- Kelvin-Helmholtz contraction and energy
- Ignited central protostar
- Shocks
- ...

Molecular Hydrogen (H₂)

- Since H₂ consists of 2 identical atoms, it has no electric dipole moment and rotationally excited H₂ has to radiate via energetically higher quadrupole transitions with excitation temperatures > 500 K.
--> cold clouds have to be observed other ways, e.g., CO

- H₂ can be detected in hot environment.
Rotational energy:

Classical mechanics: $E_{\text{rot}} = J^2/2I$

(J: Angular momentum; I: Moment of inertia)

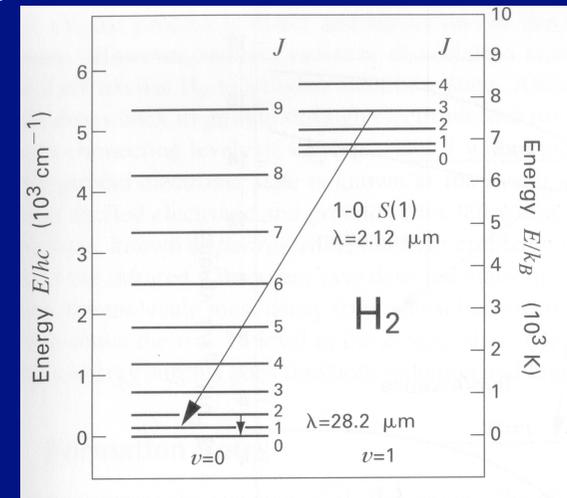
Quantum-mechanical counterpart: $E_{\text{rot}} = \frac{h^2}{2I} \times J(J+1)$
 $= BhI \times J(J+1)$

(J: rotational quantum number; B: rotational constant)

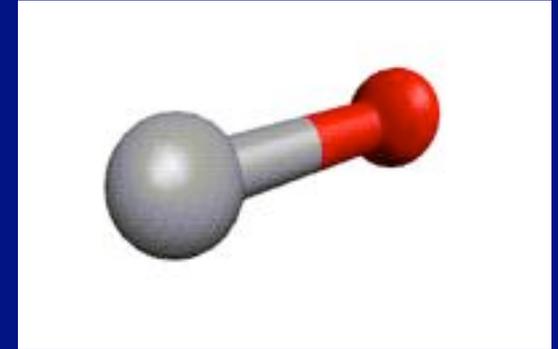
- Small moment of inertia --> large spread of energy levels

Allowed quadrupole transitions $\Delta J = 2$

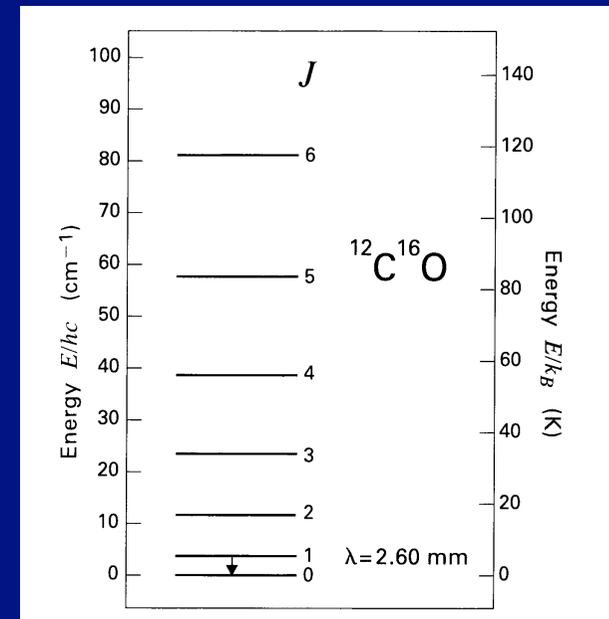
--> lowest rotational transition J=2-0 has energy change of 510 K



Carbon monoxide (CO)

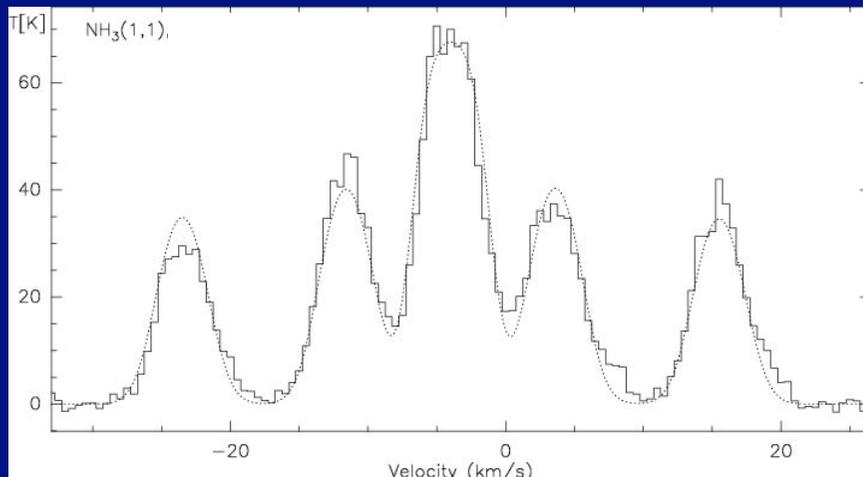
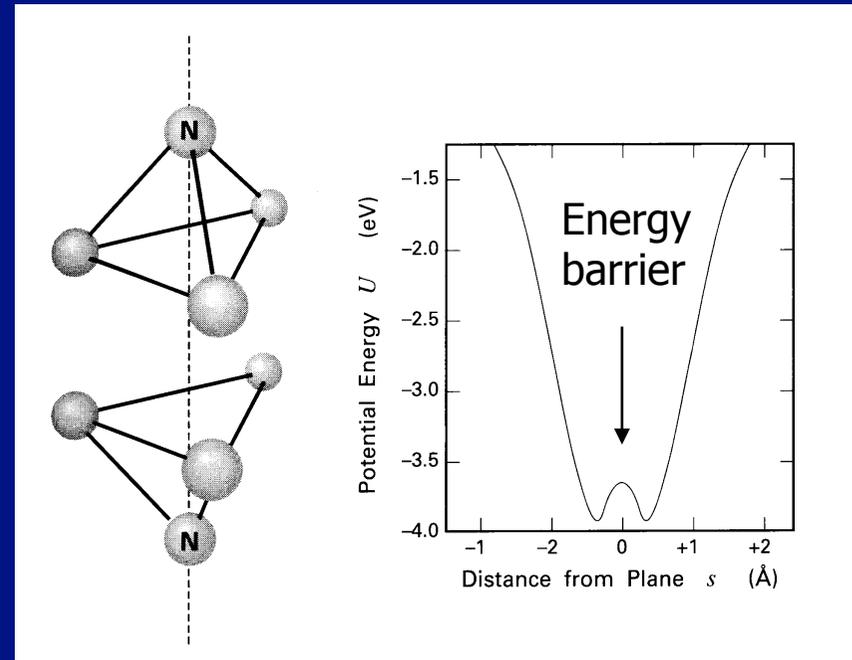
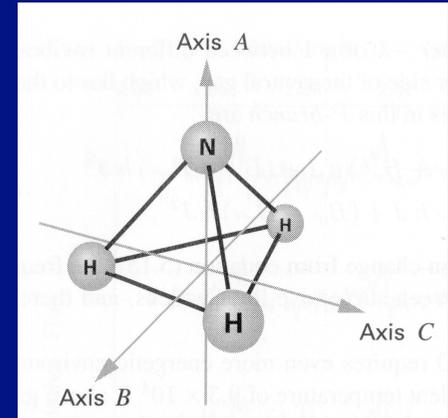


- Forms through gas phase reactions.
- Strong binding energy of 11.1 eV helps to prevent much further destruction (self-shielding).
- Has permanent dipole moment --> strong emission at (sub)mm wavelengths.
- Larger moment of inertia than H₂.
--> more closely spaced rotational ladder,
J=1 level at 4.8×10^{-4} eV or 5.5K above ground
- In molecular clouds excitation mainly via collisions with H₂.
- Critical density for thermodynamic equilibrium with H₂ $n_{\text{crit}} = A/\gamma \sim 3 \times 10^3 \text{cm}^{-3}$.
(A: Einstein A coefficient; γ : collision rate with H₂)
- The level population follows a Boltzmann-law:
$$n_{J+1}/n_J = g_{J+1}/g_J \exp(-\Delta E/k_B T_{\text{ex}})$$
 (for CO, the statistical weights $g_J = 2J + 1$)
The excitation temperature T_{ex} is a measure for the level populations and equals the kinetic temperature T_{kin} if the densities are $> n_{\text{crit}}$.



Ammonia (NH₃)

- Formed through gas-phase reactions.
- Symmetric-top molecule
- $E_{\text{rot}} = J_A^2/2I_A + J_B^2/2I_B + J_C^2/2I_C$
 - > However, useful transitions only at very high freq.
- Most useful transitions are the inversion transitions around 25GHz.
 - > tunneling energy barrier
- Additional effects (non-spherical charge distribution, quadr. mom., magn. interaction between spins) causes further hyperfine splitting.



Topics today

- Molecules and depletion
- Heating and cooling
- Radiation transfer and column density determination
- Line profiles
- Applications

Heating processes

UV radiation from stars

Energy injection from supernovae

Energy injection from outflows/jets

Cosmic rays interaction with HI and H₂

(consist mainly of relativistic protons accelerated within magnetized shocks produced by supernova-remnant--molecular cloud interactions)

$p^+ + H_2 \rightarrow H_2^+ + e^- + p^+$ (dissociation: ions also important for ion-molecule chemistry)

Interstellar radiation (diffuse field permeating interstellar space)

Mainly dissociates carbon (lower ionization potential than H₂)

$C + h\nu \rightarrow C^+ + e^-$ The electron then disperses energy to surrounding atoms by collisions.

Photoelectric heating: - Heats grains which re-radiate in infrared regime
- UV photons eject e⁻ from dust and these e⁻ heat surrounding gas via collisions

Cooling processes

Major constituents H & H₂ have no dipole moment and hence cannot effectively cool in quiescent molecular cloud. Other coolants required.

--> Hydrogen collides with ambient atoms/molecules/grains exciting them. The cooling is then done by these secondary constituents.

O + H --> O + H + hν collisional excitation (FIR)

C⁺ + H --> C⁺ + H + hν fine structure excitation (FIR)

CO + H₂ --> CO + H₂ + hν rotational excitation (radio/(sub)mm)

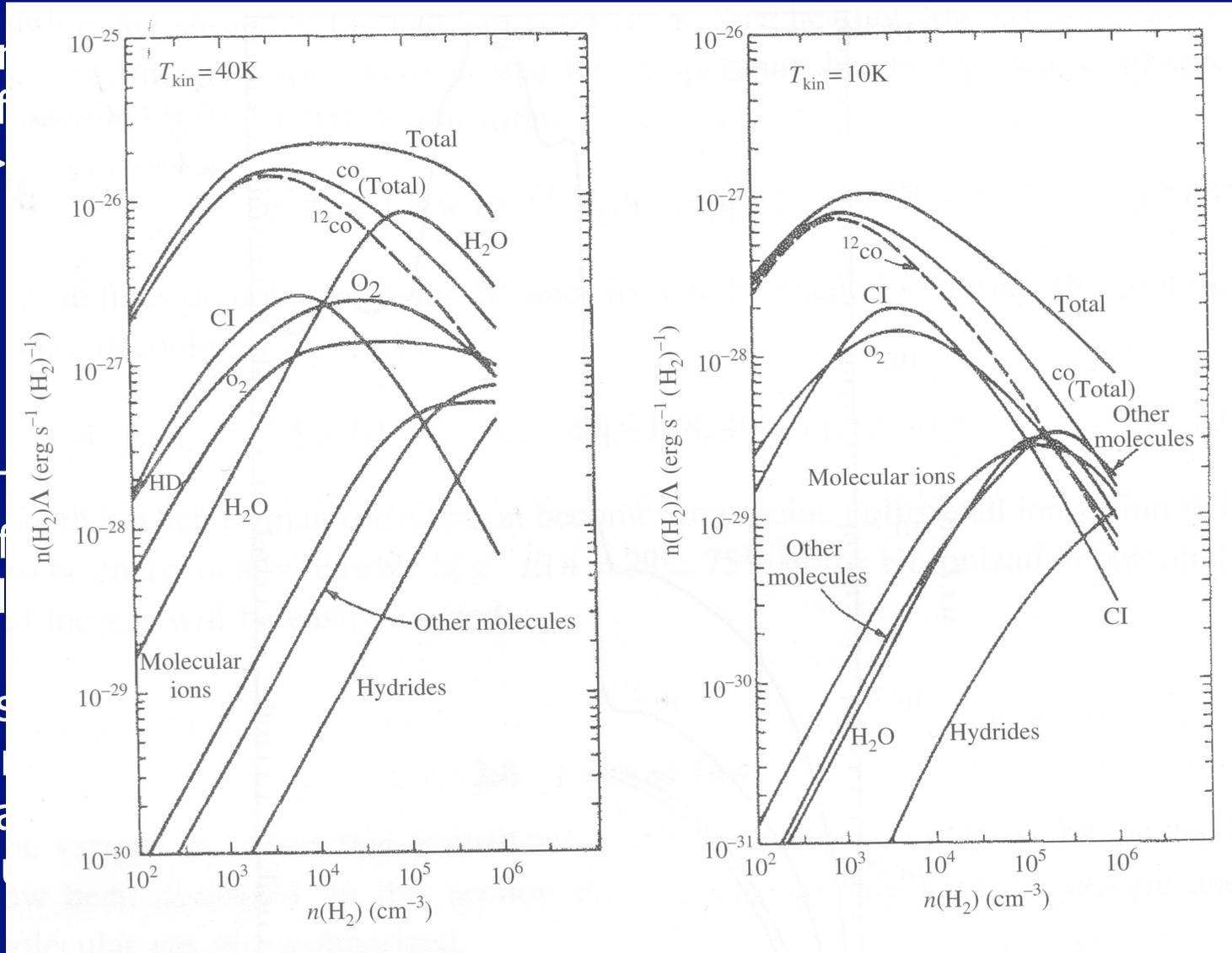
At higher densities other molecules come into play, e.g., H₂O.

The low-J CO lines are mostly optically thick, the energy diffuses from region to region and escapes from cloud surface. Higher J lines cool directly. CO is the most effective coolant in molecular clouds.

Collisions with gas atoms/molecules cause lattice vibrations on grain surfaces, that decay through the emission of infrared photons (since grains are also heated by radiation gas and dust temperature are usually not equal).

Cooling processes

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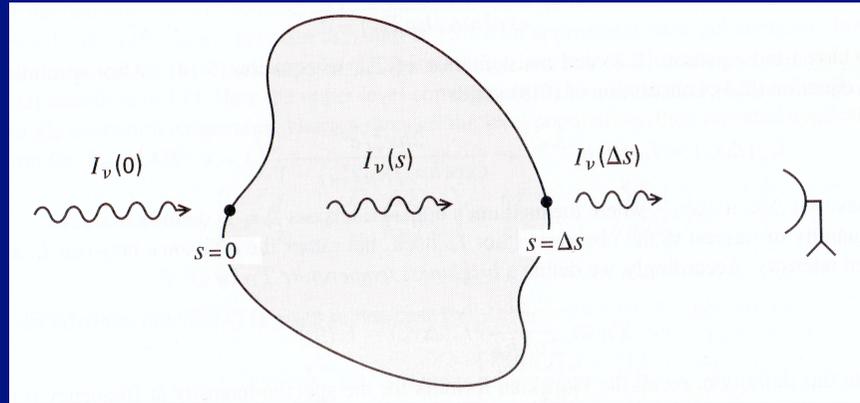


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- Heating and cooling
- **Radiation transfer and column density determination**
- Line profiles
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Radiation transfer I



$$dI_\nu = -\kappa_\nu I_\nu ds + \varepsilon_\nu ds$$

with the opacity

$$d\tau_\nu = -\kappa_\nu ds$$

and the source function

$$S_\nu = \varepsilon_\nu / \kappa_\nu$$

$$\Rightarrow dI_\nu / d\tau_\nu = I_\nu - S_\nu$$

Assuming a spatially constant source function \rightarrow radiation transfer equation

$$\Rightarrow I_\nu = S_\nu (1 - e^{-\tau_\nu}) + I_{\nu,0} e^{-\tau_\nu}$$

Radiation transfer II

The excitation temperature T_{ex} is defined via a Boltzmann distribution as

$$n_j/n_{j-1} = g_j/g_{j-1} \exp(-h\nu/kT_{\text{ex}})$$

with n_j and g_j the number density and statistical weights.

In case of rotational transitions

$$g_j = 2J + 1$$

In thermal equilibrium

$$T_{\text{ex}} = T_{\text{kin}}$$

In a uniform molecular cloud the source function S_ν equals Planck function

$$S_\nu = B_\nu(T_{\text{ex}}) = 2h\nu^3/c^2 (\exp(h\nu/kT_{\text{ex}}) - 1)^{-1}$$

And the radiation transfer equation

$$\Rightarrow I_\nu = B_\nu(T_{\text{ex}}) (1 - e^{-\tau_\nu}) + I_{\nu,0} e^{-\tau_\nu}$$

In the Rayleigh-Jeans limits ($h\nu \ll kT$) B equals

$$B = 2k\nu^2/c^2 T \quad (\text{def. } \rightarrow T = c^2/(2k\nu^2) I_\nu)$$

And the radiation transfer equation using now the radiation temperature is

$$T_r = J_\nu(T_{\text{ex}}) (1 - e^{-\tau_\nu}) + J_{\nu,0}(T_{\text{bg}}) e^{-\tau_\nu}$$

With

$$J_\nu = h\nu/k (\exp(h\nu/kT) - 1)^{-1}$$

Subtracting further the background radiation

$$T_r = (J_\nu(T_{\text{ex}}) - J_{\nu,0}(T_{\text{bg}})) (1 - e^{-\tau_\nu})$$

Molecular column densities

To derive molecular column densities, 3 quantities are important:

- 1) Intensity T of the line
- 2) Optical depth τ of the line (observe isotopologues or hyperfine structure)
- 3) Partition function Q

The optical depth τ of a molecular transition can be expressed like

$$\tau = c^2/8\pi\nu^2 A_{ul}N_u (\exp(h\nu/kT) - 1) \phi$$

With the Einstein A_{ul} coefficient

$$A_{ul} = 64\pi^4\nu^3/(3c^3h) \mu^2 J_u/(2J_u-1)$$

And the line form function ϕ

$$\phi = c/\nu 2\text{sqrt}(\ln 2)/(\text{sqrt}(\pi)\Delta\nu)$$

Using furthermore the radiation transfer eq. ignoring the background

$$T = J_\nu (T_{ex}) \tau (1 - e^{-\tau})/\tau$$

And solving this for N_u , one gets

$$N_u = 3k/8\pi^3\nu 1/\mu^2 (2J_u-1)/J_u \tau / (1 - e^{-\tau}) (T\Delta\nu \text{sqrt}(\pi)/(2\text{sqrt}(\ln 2)))$$

The last expression equals the integral $\int T dv$.

The column density in the upper level N_u relates to the total column density N_{tot}

$$N_{tot} = N_u/g_u Q \exp(E_u/kT)$$

For a linear molecule like CO, the partition function can be approximated to

$$Q = kT/hB.$$

However, for more complex molecules Q can become very complicated.

Conversion from CO to H₂ column densities

One classical way to derive conversion factors from CO to H₂ column densities and gas masses essentially relies on three steps:

- 1) Derive ratio between colour excess E_{B-V} and optical extinction A_V
$$A_V = 3.1 E_{B-V} \quad (\text{Savage and Mathis, 1979})$$
 - 2) The ratio $N(\text{H}_2)/E_{B-V}$: One can measure the H₂ column density, e.g., directly from UV Absorption lines.
 - 3) The ratio $N(\text{CO})/A_V$: In regions of molecular gas emission, one can estimate A_V by star counts in the Infrared regime
- ⇒ Combining these three ratios, the CO observations can directly be converted to H₂ column densities. Assumptions about the 3D cloud geometry allow further estimates about the cloud masses and average densities.

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Line broadening

Natural line broadening: Disturbance of molecule by zero-point vibrations of electromagnetic field (or from thermal electromagnetic field)

$$dv = 32\pi^3\nu^3 \mu^2 / (3hc^3) \quad (\mu: \text{Dipole moment})$$

For CO(1-0) at 20K $\rightarrow dv \sim 3.5 \times 10^{-8}$ or $dv \sim 9 \times 10^{-14}$ km/s
 \rightarrow Negligible!

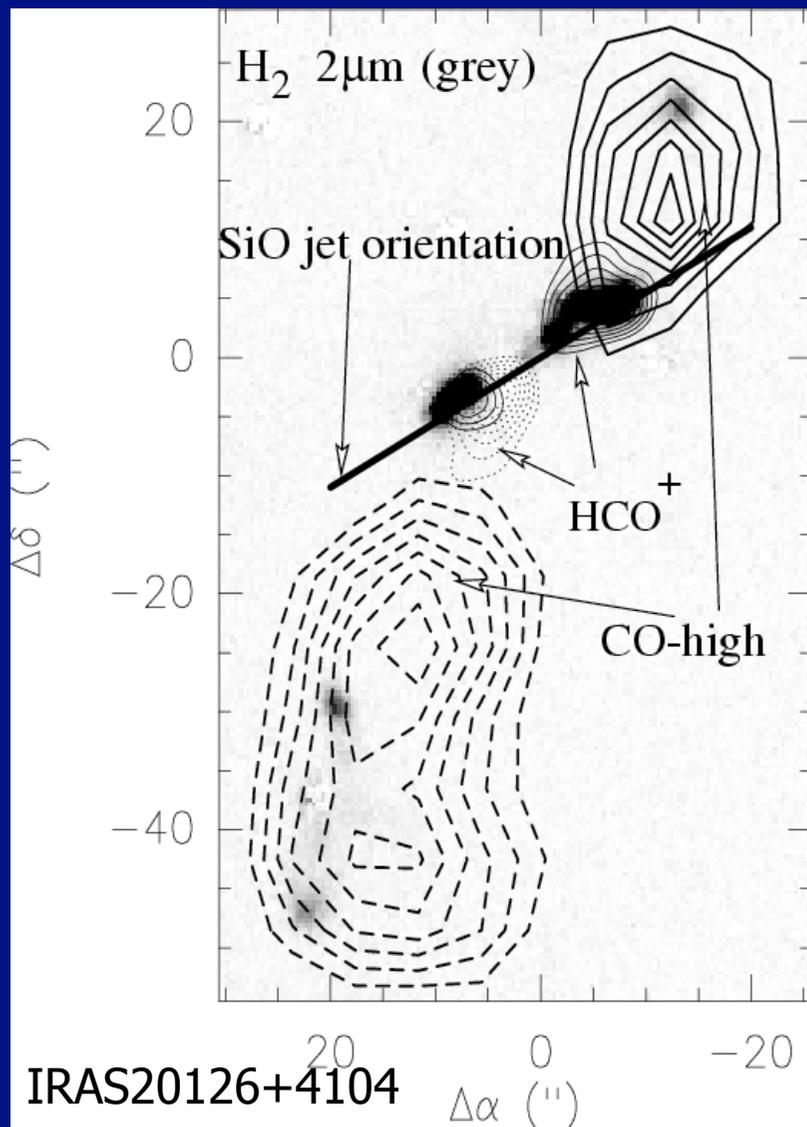
Pressure broadening: Arises from collisions between molecules. Complex quantum-mechanical problem for intermolecular forces. At densities of star-forming regions negligible.

Thermal line broadening: Thermal motions of gas cause doppler broadening:

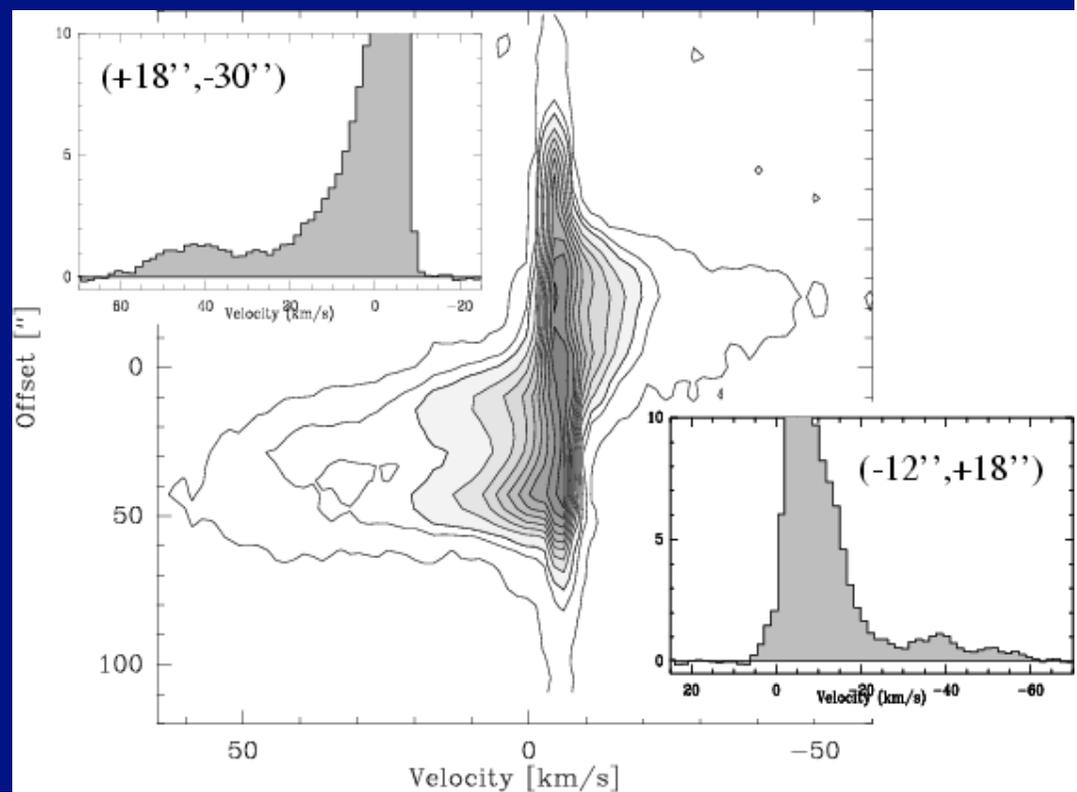
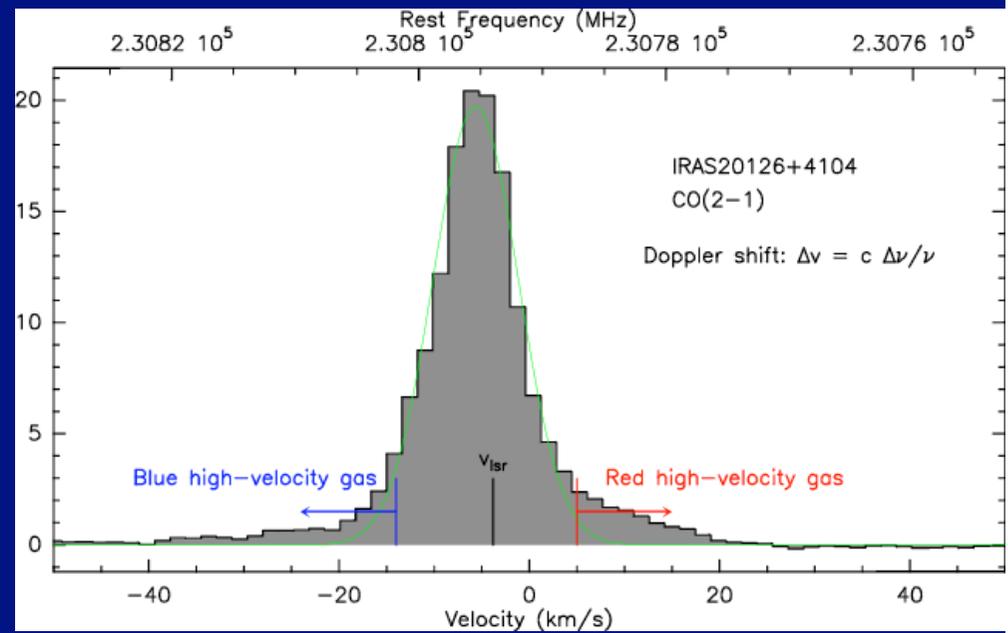
$$dv = \text{sqrt}(8\ln 2 \text{ kT} / m_{\text{mol}})$$
$$dv(\text{NH}_3 @ 30\text{K}) \sim 0.28 \text{ km/s}$$

Other physical effects: Line broadening due to outflow motions, rotation ...

Molecular outflows



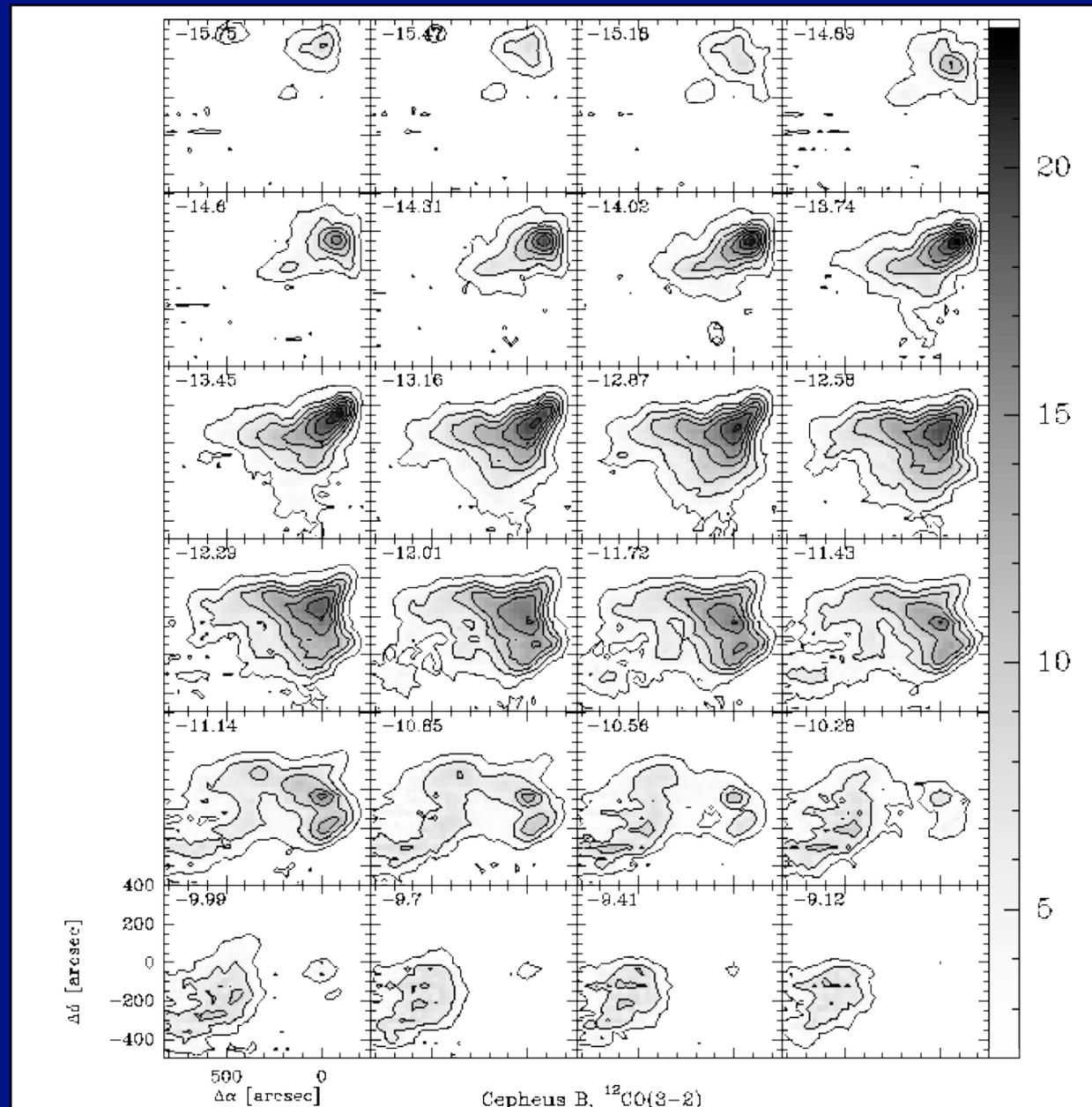
Lebron et al. 2006



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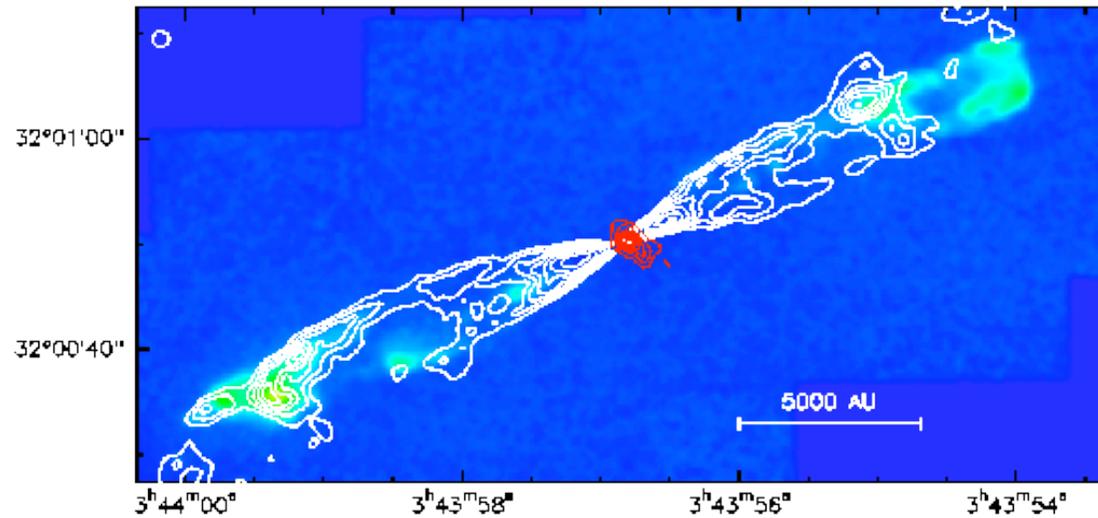
Applications II: Velocity structure of molecular clouds



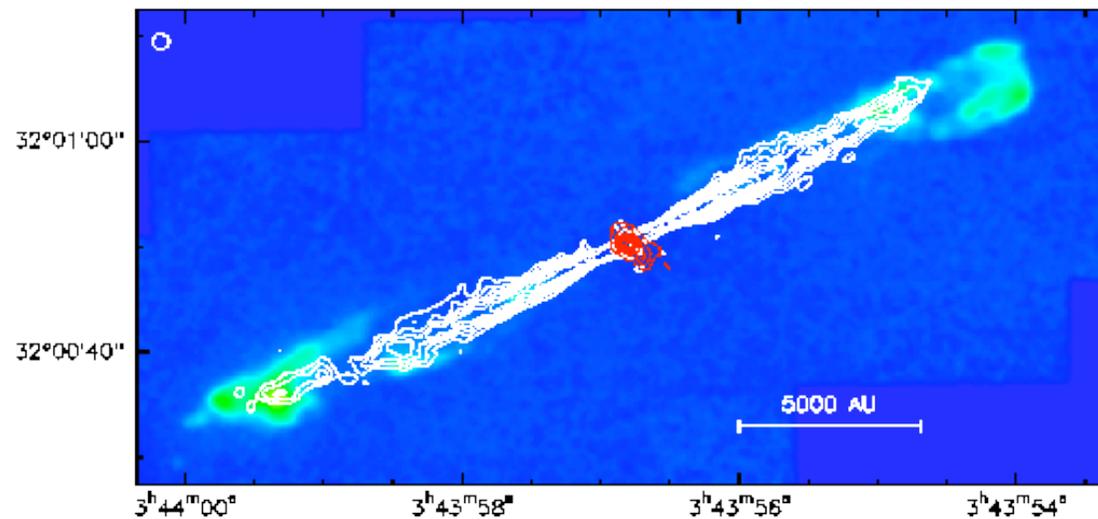
Applications III: Molecular outflows

HH211, Gueth et al. 1999

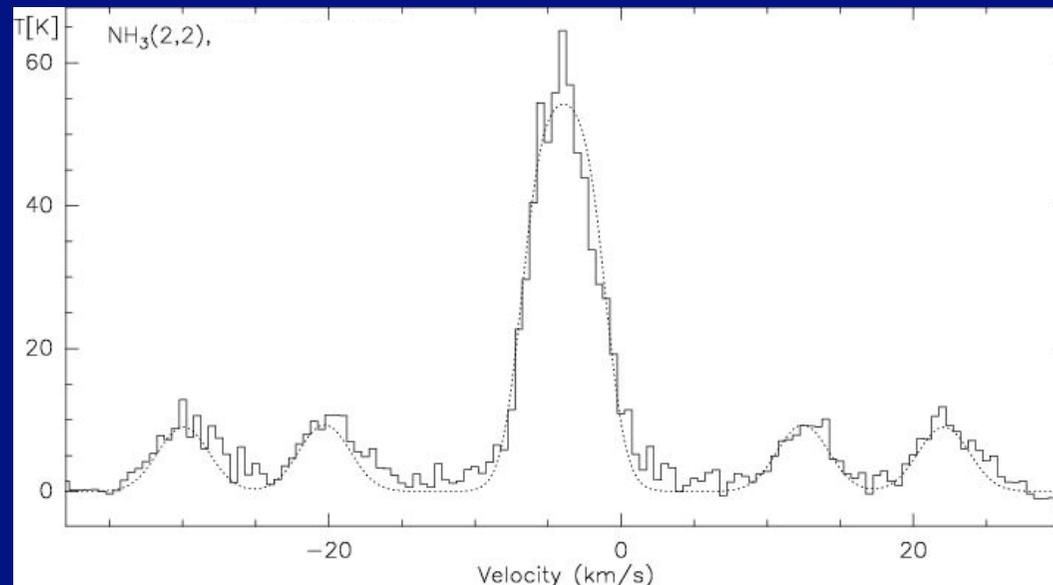
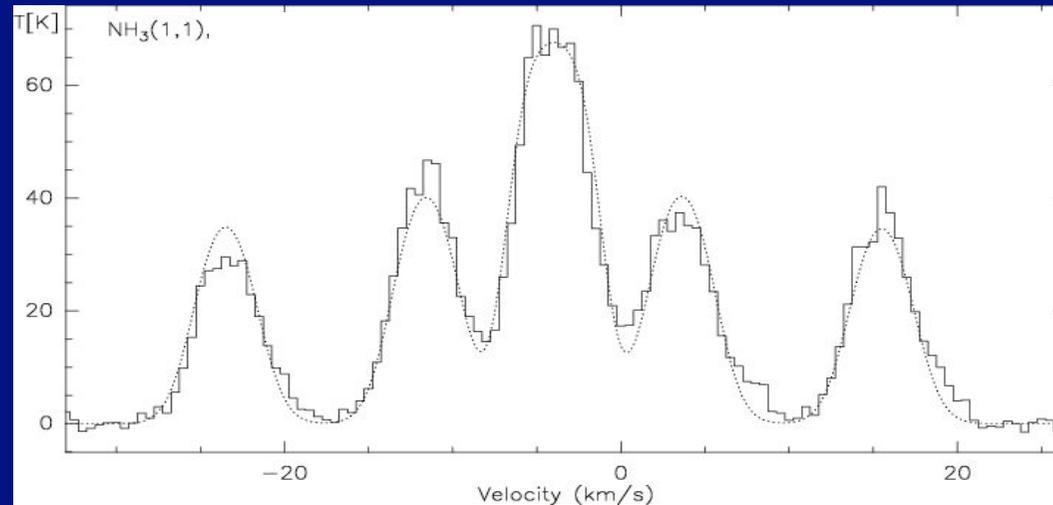
H₂ 2.12 μm (colors) + CO J=2-1 v<10 km/s (white) + continuum 1.3 mm (red)



H₂ 2.12 μm (colors) + CO J=2-1 v>10 km/s (white) + continuum 1.3 mm (red)



Applications IV: Temperature estimates from NH₃



$$T_{\text{kin}} = T_{\text{rot}} (\tau_{11} \tau_{22} T_{11} T_{22})$$

Summary

- Main tools: Spectral line emission and thermal emission and extinction from dust (more on dust next week)
- Molecules interesting for themselves and chemistry
- However, also extremely useful to trace physical processes.
- Molecules deplete on grains at low temperatures
- Discussed main cooling and heating processes
- Discussed basic line radiation transfer and column density determination
- What causes line broadening?
- A few applications

Next week more about dust!

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