Sternentstehung - Star Formation Winter term 2022/2023 Henrik Beuther, Thomas Henning & Jonathan Henshaw 18.10 Today: Introduction & Overview (Beuther) 25.10 Physical processes I (Beuther) 08.11 Physcial processes II (Beuther) (Henshaw) 15.11 Molecular clouds as birth places of stars 22.11 Molecular clouds (cont.), Jeans Analysis (Henshaw) 29.11 Collapse models I (Beuther) 06.12 Collapse models II (Henning) 13.12 Protostellar evolution (Beuther) 20.12 Pre-main sequence evolution & outflows/jets (Beuther) (Henning) 10.01 Accretion disks I (Henning) 17.01 Accretion disks II 24.01 High-mass star formation, clusters and the IMF (Henshaw) 31.01 Extragalactic star formation (Henning) 07.02 Planetarium@HdA, outlook, questions 13.02 Examination week, no star formation lecture Book: Stahler & Palla: The Formation of Stars, Wileys

More Information and the current lecture files: http://www.mpia.de/homes/beuther/lecture_ws2223.html beuther@mpia.de, henning@mpia.de , henshaw@mpia.de

Topics today

- The ISM, molecules and depletion

- Heating and cooling

Radiation transfer and column density determination

The cosmic cycle



http://www.astro.uni-koeln.de

Properties of Molecular Clouds

| Туре | n [cm ⁻³] | Size [pc] | Т [K] | Mass [M _{sun}] |
|-----------------------|--------------------------|--------------|----------|-----------------------------|
| Giant Molecular Cloud | 10 ² | 50 | 15 | 10 ⁵ |
| Dark Cloud Complex | 5x10 ² | 10 | 10 | 104 |
| Individual Dark Cloud | 10 ³ | 2 | 10 | 30 |
| Dense low-mass cores | 104 | 0.1 | 10 | 10 |
| Dense high-mass cores | >105 | 0.1-1 | 10-30 | 100-10000 |



Neutral and ionized medium

atomic hydrogen

http://adc.gsfc.nasa.gov

S. LANG A.

radio continuum (2.5 GHz)

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

The Interstellar Medium I

Atomic Hydrogen



<u>21cm line:</u> electron spin S flip from parallel (F=1) to antiparallel (F=0) compared to the Proton spin I.

atomic hydrogen

http://adc.gsfc.nasa.gov

 $\Delta E = 5.9 \times 10^{-5} \, eV$

The Interstellar Medium I



Energy, E

The Ionized gas

Ionized gas

http://adc.gsfc.nasa.gov

radio continuum (2.5 GHz)

- Hydrogen recombination lines from optical to cm wavelengths
- Emission lines from heavier elements --> derive atomic abundances

He/H

C/H

N/H

- Free-free emission between e⁻ and H⁺



0.1

3.4x10⁻⁴

6.8x10⁻⁵

The Molecular ISM

Molecular Hydrogen



http://adc.gsfc.nasa.gov



Formaldehyde H₂CO

Cyanoacetyline HC₃N

molecular hydrogen



Excitation mechanisms:

- Rotation
- Vibration
- Electronic transitions

--> usually cm and (sub)mm wavelengths

- --> usually submm to FIR wavelengths
- --> usually MIR to optical wavelengths

Molecular ISM Basics

History:

- Late 1930s: Detection of CH, CH⁺ and CN in diffuse clouds by absorption of optical light from background stars
- 1960s: Detection of OH, NH₃ and H₂O at radio wavelength
- 1970: CO

Formation of molecules is an energy problem: Two atoms approach each other with positive total energy → rebound if no energy can be given away

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
 --> probalility of such radiative association low as well

Molecular ISM Basics

- Ion-molecule or ion-atom reactions can solve energy problem
- Neutral-neutral reactions on dust grain surfaces (catalytic) important
- Ion induces dipole moment in atom or molecule
 --> creates electrostatic attraction between the two.
 --> effective cross section increases over geometric values
- At low temperatures such reactions account for large fraction of molecules.
- However, not enough ions to account for large H_2 abundances --> grain surface chemistry important

Simple molecules like CO or CS → ion-molecule chemistry,
 More complex molecules → grain surface chemistry important

Molecular ISM Basics



- However, not enough ions to account for large H₂ abundances --> grain surface chemistry important

- Simple molecules like CO or CS \rightarrow ion-molecule chemistry,

- More complex molecules \rightarrow grain surface chemistry important

Molecules in Space

| 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 atoms |
|--|--|--|--|--|--|--|--|---|------|----------|----------|
| H2 AIF AICI C2 CH + CN CO C9 CSi HCI NO NS NaCI OH NS S0 S0 SiS SS HF FeO(?) | C3 C2H C2O C2S CH2 HCN HCO HCO+ HCO+ HCS+ HOC+ H2O H2S HNC HNO MgNC N2H+ N2O NaCN OCS SO2 c-SiC2 CO2 NH2 H3+ SiCN AINC | c-C3H I-C3H C3N C3O C3S C2H2 CH2D+? HCCN HCNH+ HNCO HNCS HOCO+ H2CO H2CN H2CS H3O+ NH3 SiC3 C4 | C5 C4H C4Si I-C3H2 c-C3H2 CH2CN CH4 HC3N HC2NC HCOOH H2CHN H2C2O H2NCN HNC3 SiH4 H2COH+ | C5H I-H2C4 C2H4 CH3CN CH3NC CH3OH CH3SH HC3NH+ HC2CHO NH2CHO C5N HC4N | C6H CH2CHCN CH3C2H HC5N HCOCH3 NH2CH3 c-C2H4O CH2CHOH | CH3C3N HCOOCH3 CH3COOH? C7H H2C6 CH2OHCHO CH2CHCHO | СН3С4Н СН3СН2СN (СН3)2О СН3СН2ОН НС7N С8Н | CH3C5N? (CH3)2CO NH2CH2COOH? CH3CH2CHO | HC9N | CH3OC2H5 | HC11N |

About 270 detected interstellar molecules as of October 2022 (<u>www.cdms.de</u>). 73 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 ∏ _{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_2O | 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 | 1×10 ⁵ | Dense gas/temperature probe |
| CH ₃ CN | 19-18 | 2x10 ⁻⁸ | 2x10 ⁷ | Temperature probe in Hot Cores |

Crit. Dens.: $n_{crit} \sim A/\gamma$

Basics IV

Depletion of molecules on dust grains

In molecule's ref. frame, grains are moving at v_{therm} relative to molecules

 $E = 1/2 \text{ mv}_{\text{therm}}^2 = 3/2 \text{ k}_{\text{b}} \text{T} => \text{v}_{\text{therm}} = (3\text{k}_{\text{b}}\text{T/m})^{1/2}$

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

Probability of molecule in volume V to be struck by grain in time $\Delta t P(\Delta t) = n(\pi a^2)v_{therm}\Delta t /V$

For example CS: $v_{therm} \sim 5x10^3$ cm s⁻¹ at 10K, $n_H \sim 10^4$ cm⁻³, $\Sigma \sim 10^{21}$ cm² $t_{coll} \sim 6x10^5$ yr

Depletion time-scale very short --> mechanisms for re-injecting molecules from grains important

Depletion example

1.2 mm Dust Continuum

C¹⁸O





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)

- H_2 consists of 2 identical atoms \rightarrow no electric dipole moment

- Rotationally excited H₂ has allowed quadrupole transitions $\Delta J = 2$ \rightarrow lowest rotational transition J=2-0 has energy change of 510 K

 Rotational energy for H₂: Classical mechanics: E_{rot} = J²/2I (J: Angular momentum; I: Moment of inertia)

→ Small moment of inertia (I=mr²)
→ large spread of energy levels



 \rightarrow Cold clouds have to be observed other ways, e.g., CO

Carbon monxide (CO)

- Forms through gas phase reactions.
- Strong binding energy of 11.1 eV
 - \rightarrow prevents much further destruction (self-shielding).
- Permanent dipole moment \rightarrow strong emission at (sub)mm wavelengths.
- Larger moment of inertia than H₂.
 → more closely spaced rotational ladder, J=1 level at 4.8x10⁻⁴eV or 5.5K above ground
- In molecular clouds excitation mainly via collisions with H₂.
- Critical density for thermodynamic equilibrium with H₂ n_{crit} = A/γ ~ 3x10³cm⁻³. (A: Einstein A coefficient; γ: collision rate with H₂)







http://www.cdms.de

Topics today

- The ISM, molecules and depletion

- Heating and cooling

Radiation transfer and column density determination

Heating processes

UV radiation from stars

Energy injection from supernovae

Energy injection from outflows/jets

 Cosmic rays interact 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 \rightarrow ion-molecule chemistry)

 Interstellar radiation (diffuse field permeating interstellar space) Mainly dissociates carbon (lower ionization potential than H₂)

 $C + hv \rightarrow C^+ + e^-$ Electron disperses energy to ISM 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

- H & H₂ no dipole moment \rightarrow no efficient coolant in cold molecular cloud \rightarrow other coolants needed

--> Hydrogen collides with ambient atoms/molecules/grains \rightarrow Cooling via these secondary constituents.

O + H --> O + H + hv collisional excitation (FIR) $C^+ + H \rightarrow C^+ + H + hv$ fine structure excitation (FIR) $CO + H_2 --> CO + H_2 + h_V$ rotational excitation (radio/(sub)mm) At higher densities other molecules come into play, e.g., H_2O .

 \rightarrow CO 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.

 \rightarrow dust very efficient coolant

Cooling processes



Topics today

- The ISM, molecules and depletion

- Heating and cooling

Radiation transfer and column density determination

Radiation transfer I



 $dI_{v} = -\kappa_{v}I_{v,0}ds + \varepsilon_{v}ds$

with the opacity $d\tau_v = -\kappa_v ds$

κ: absorption coef.ε: emission coef.

and the source function $S_v = \epsilon_v / \kappa_v$

 $\Rightarrow dI_{\rm v} / \ d\tau_{\rm v} = I_{\rm v,0} - S_{\rm v}$

Assuming a spatially constant source function \rightarrow radiation transfer equation

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

Radiation transfer II

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

 $n_{\rm J}/n_{\rm J-1} = g_{\rm J}/g_{\rm J-1} \exp(-h_{\rm V}/kT_{\rm ex})$

with n_{J} and g_{J} the number density and statistical weights.

In case of rotational transitions

 $g_{J} = 2J + 1$

J: rot. quantum number

In thermal equilibrium

 $T_{ex} = T_{kin}$

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

 $S_v = B_v (T_{ex}) = 2hv^3/c^2 (exp(hv/kT_{ex}) - 1)^{-1}$

Radiation transfer III

Then the radiation transfer equation

 $\Rightarrow I_{v} = \mathsf{B}_{v} (\mathsf{T}_{ex}) (1 - e^{-\tau(v)}) + I_{v,0} e^{-\tau(v)}$

In the Rayleigh-Jeans limits (hv < < kT) B equals

 $B = (2kv^2/c^2)T$ (def. $\rightarrow T = c^2/(2kv^2)I_v$)

And the radiation transfer equation using now the radiation temperature is

$$T_{r} = J_{v} (T_{ex}) (1 - e^{-\tau(v)}) + J_{v,0} (T_{bg}) e^{-\tau(v)}$$

with

 $J_v = hv/k (exp(hv/kT) - 1)^{-1}$

Molecular column densities I

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 $\boldsymbol{\tau}$ of a molecular transition can be expressed like

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

with the Einstein A_{ul} coefficient

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

and the line form function $\boldsymbol{\phi}$

 $\phi = c/v 2 \operatorname{sqrt}(\ln 2) / (\operatorname{sqrt}(\pi) \Delta v)$

Molecular column densities II

Using furthermore the radiation transfer eq. ignoring the background

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

And solving τ -equation for N_u, one gets

 $N_{u} = \frac{3k}{8\pi^{3}\nu} \frac{1}{\mu^{2}} (2J_{u}-1)/J_{u} \tau / (1 - e^{-\tau}) (T\Delta\nu \operatorname{sqrt}(\pi)/(2\operatorname{sqrt}(\ln 2)))$

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

 $\rightarrow N_u \sim \tau / (1 - e^{-\tau}) \int T dv$

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

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

For linear molecule like CO, partition function Q approximated: Q = kT/hB. (B: rotational constant)

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

Conversion from CO to H₂ column densities

Classical way to derive conversion factors from CO to H₂ column densities:

- 1) Derive ratio between colour excess E_{B-V} and optical extinction $A_v = 3.1 E_{B-V}$ (Savage and Mathis, 1979)
- 2) The ratio $N(H_2)/E_{B-V}$: One can measure the H₂ column density, e.g., directly from UV Absorption lines.
- 3) The ratio $N(CO)/A_v$: In regions of molecular gas emission, one can estimate A_v by star counts in the Infrared regime
- \Rightarrow Combining these three ratios: CO \rightarrow H₂ column densities.

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

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Heidelberg Joint Astronomical Colloquium Winter Semester 2022 — Tuesday October 25th, 16:00 Main Lecture Theatre, Philosophenweg 12 Sarbani Basu (Yale University, USA): The Sun as a variable Star



Those unable to attend the colloquium in person are invited to participate online through Zoom. More information is given on HePhySTO: <u>https://www.physik.uni-heidelberg.de/hephysto/</u>