

Review of radiative transfer and thermodynamics

- Radiative transfer
 - Process
 - Quantities ($I_\nu, j_\nu, \alpha_\nu, S_\nu, \tau_\nu$)
 - Transport equation
 - Formal solution
 - Quantities derived from I_ν
- Local thermodynamical equilibrium (LTE)
 - Planck function
 - RT equation
 - Radio astronomy:
R-J approximation, brightness temperature
- Measurement of interstellar lines
 - Brightness temp. \rightarrow cloud temperature
 - Equivalent width \rightarrow column density
- State of the gas
 - Speed distribution
 - Equation of state

Interstellar spectral lines

- Atoms and ions
 - Hydrogen 21 cm
 - Recombination lines
 - IR fine structure lines
 - Absorption lines
- Molecules
 - Classification
 - Diatomic molecules:
 - * Energy levels
 - * Spectra

Spectra of diatomic molecules

- Electronic transitions
 far-UV: H_2 , CO, OH
 near-UV: CH, CH^+ , CN
 near-IR: C_2 , CN
- Vibrational transitions
 Energy levels: $E_v = (v + \frac{1}{2})h\nu_e$ (approximation)
 $v = 0, 1, 2, \dots$ vibrational quantum number;
 $\nu_e \dots$ vibrational frequency
 Selection rules: $\Delta v = \pm 1$ (fundamental transition)
 $\rightarrow \Delta E = h\nu_e \rightarrow$ one line at $\nu_e \approx 10\mu\text{m} \rightarrow$ infrared
 $\Delta v = \pm 2, \pm 3, \dots$ overtones
- Rotational transitions
 Energy levels: $E_J = BJ(J + 1)$
 $J = 0, 1, 2, \dots$ rotational quantum number
 $B = \frac{h^2}{8\pi^2\overline{m}r^2}$; $\overline{m} = \frac{m_A m_B}{m_A + m_B}$ reduced mass
 Selection rules: $\Delta J = \pm 1$,
 no lines for symmetric molecules (e.g. H_2 , O_2 , CH_4)
 $\Delta E = h\nu = 2B(J + 1)$
 \rightarrow equidistant lines with separation $2B$
 $B \propto \frac{1}{\overline{m}}$
 \rightarrow lines from lighter molecules at higher frequencies (shorter wavelengths) with larger separation between lines
 lines are at mm wavelengths (microwave)
 Example: CO $\lambda_{1 \rightarrow 0} = 2.6 \text{ mm}$,
 $\lambda_{4 \rightarrow 3} = \lambda_{1 \rightarrow 0}/4 = 0.65 \text{ mm}$, $\lambda_{10 \rightarrow 9} = 0.26 \text{ mm}$

- Vibration-rotation transitions (within one electronic state)

Combined energy levels and selection rules:

$$E_{vJ} = (v + \frac{1}{2})h\nu_e + B_v J(J+1)$$

$$\Delta v = \pm 1, \Delta J = \pm 1$$

$$v = 0 \rightarrow v = 1: B_1 < B_0 \text{ because } r_1 > r_0 \text{ and } B \propto \frac{1}{r^2}$$

$$* J \rightarrow J+1: \Delta E = h\nu_e + 2B_1 + (3B_1 - B_0)J + (B_1 - B_0)J^2$$

→ for each J a line at $\nu > \nu_e$ → “R branch”

frequency intervals $\Delta\nu$ decrease with J until they become 0 at a maximum ν , then negative with $|\Delta\nu|$ increasing (→ “band head” at large J , e.g. $J = 108$ for CO, → usually not visible)

$$* J \rightarrow J-1: \Delta E = h\nu_e - (B_1 + B_0)J + (B_1 - B_0)J^2$$

→ for each J a line at $\nu < \nu_e$ → “P branch”

frequency intervals $|\Delta\nu|$ increase with J

- Intensity of rotational transitions determined by

1. Transition dipole moment

2. Populations of levels

calculated from Boltzmann statistics in LTE:

$$P_J = \frac{N_J}{N} = (2J+1) \exp(-BJ(J+1)/kT) / (kT/(sB))$$

$s \dots 2$ or 1 for symmetric or nonsymmetric molecules

(assumption: only ground vibrational and electronic states are populated)

$$J \text{ at maximum population } (\frac{dN_J}{dJ} = 0): J_{\max} \propto \sqrt{T}$$

→ cloud temperatures can be measured from relative intensities of rotational lines

Cooling in the ISM

- Criteria for efficient cooling

1. High abundances of colliding particles

Atoms: H, C, N, O; e^- ;

Molecules: H_2 (electric quadrupole transitions), CO at ≈ 10 K

2. $E_{\text{kin}} \approx E_{\text{exc}}$

Transition	$\Delta E/k$
C II ($^2P_{1/2} \rightarrow ^2P_{3/2}$)	92 K
Si II ($^2P_{1/2} \rightarrow ^2P_{3/2}$)	413 K
O I ($^3P_2 \rightarrow ^3P_{1,0}$)	228 K
	326 K
H_2 ($\Delta J = \pm 2$)	510 K
CO ($J = 0 \rightarrow 1$)	6 K

3. Large collisional excitation cross-section \rightarrow ions + e^-

4. $A_{\text{ul}} > C_{\text{ul}}$ (strong transitions)

5. $\tau < 1 \rightarrow$ balance between 1. and 4.

not satisfied for CO in dense clouds

\rightarrow OH, H_2O , ^{13}CO , C^{18}O important coolants

- Cooling rate Λ

loss of thermal energy per unit volume and per second

$$\Lambda_{\text{C}^+} \propto n(\text{C}^+) \cdot n(e^-) \cdot \frac{1}{\sqrt{T}}$$

- Cooling time t_c

time necessary for a significant change of temperature:

ratio between internal energy of the gas per unit volume

and cooling rate

Heating processes

1. Starlight

- Photoionization of C, Si, Fe
mean $\Delta Q \approx \mathbf{2.1}$ eV per ionization
- Photodissociation of H_2
excitation into state with larger internuclear separation than ground state
mean $\Delta Q \approx \mathbf{0.4}$ eV per dissociation

2. Cosmic rays ($p + e^-$ with $E \approx \text{MeV}$)

and soft X-rays ($E \approx 1 \text{ keV}$)

- Ionization of H by cosmic rays
mean $\Delta Q \approx \mathbf{3.4}$ eV per ionization
- Ionization of He by X-rays
mean $\Delta Q \approx \mathbf{6}$ eV per ionization
- Ionization of H_2 by CRs and X-rays
 $\Delta Q \approx \mathbf{15}$ eV per ionization for 2 MeV proton

3. Grains

photoelectric effect (UV photons)

mean E_{kin} of liberated electron $\approx \mathbf{5}$ eV

important in diffuse neutral clouds

4. (Magnetohydrodynamic heating

- Stellar winds
- Supernova explosions
- Expansion of HII regions
- Gravitational collapse of a cloud
- Magnetic fields

Molecule formation

- Problems for direct formation $A + B \rightarrow AB + E$
 - Short collision contact time
 - Low transition probability
 - Very low third body collision probability
- Solutions
 - Involvement of ions
 - Dust grain catalysis (e.g. H_2)
 - Indirect formation by chemical reactions $A + B \rightarrow C + D$
- Ion-molecule reactions
 - Very rapid
 - Rates independent of temperature
 - Rates similar for different molecules
- Neutral exchange reactions
 - Lower rates than ion-molecule reactions
 - Rates depend on temperature
 - Rates similar for different molecules
- Reaction networks
 - Molecules formed and destroyed by above two types of reactions occurring successively, starting from H_2
+ destroyed by UV radiation
 - Set of differential equations for the change of the number densities of all molecules with time
 - Equilibrium abundances for $\frac{dn(M)}{dt} = 0$
 - Examples: Ionization level and number density of atomic hydrogen in dense clouds, CO formation

Interstellar dust

- Evidence for dust grains
 - Element abundances in the ISM
 - Extinction of starlight
 - Polarization of starlight
 - Scattered light
 - Spectral features from dust (absorption + emission)
 - Photoluminescence
- Formation of grains (condensation)
 - IS clouds (slow)
 - Atmospheres of cool giants (fast)
 - Envelopes of protostars
- Destruction of grains
 - Evaporation
 - Sputtering in shocks (e.g. SN blast wave)
- Grain properties
 - Temperature (10 ... 50 K, time dependent for small grains)
 - Charge
 - * 1 ... 100 e, depending on temperature, from collisions
 - * positive charge from photoelectric effect
- H₂ formation on grains (DW 4.6 and essay topic)

Radiatively excited regions

- Regions of high temperatures and low densities:
Clouds of atomic H near hot stars
 - T up to 10000 K, $n \approx 10^8 - 10^9 \text{ m}^{-3}$
 - HII regions: clouds around stars
 - “diffuse ionized medium”: lower densities, not clearly associated with stars
 - Observed spectra show emission lines:
H Balmer, forbidden lines from O, N, S ions
- Hydrogen nebulae
 - Ionization of H by UV radiation from star
 - Recombination of $p + e^-$
 - Ionization equilibrium
 - Degree of ionization (x): $x \approx 1$
 - Variation of x with distance from star: well defined edge
 - Strömgren radius R_s
 - Temperature T_e from energy balance: $T_e \approx T_{\text{eff}}$
- Heavier elements in HII regions
 - Oxygen-line cooling
 - Ionization stratification
- Radio continuum spectrum
- Radio line spectrum (DW 5.4.2)
- Determination of electron temperature and density from observations