ASTR 8030/3600 Stellar Astrophysics

X-ray Sunday

SDO 4500 Angstroms: photosphere

SDO/AIA 4500 2012-08-20 12:00:07 UT

SDO 1600 Angstroms: upper photosphere

T~5x104K

SDO/AIA 1600 2012-08-20 10:58:41 UT

SDO 304 Angstroms: chromosphere

 SDO/AlA 304 2010-08-23 00:07:21 UT

SDO 171 Angstroms: quiet corona

SDO/AIA 171 2012-08-20 16:39:00 UT

SDO 211 Angstroms: active corona

T~2x106K

SDO/AIA 211 2012-08-20 16:22:13 UT

SDO 94 Angstroms: flaring regions

T~6x106K

 SDO/AIA 94 2012-08-20 16:46:14 UT

SDO: dark plasma (3/27/2012)

SDO: solar flare (4/16/2012)

SDO: coronal mass ejection (7/2/2012)

Aims of the course

- Introduce the equations needed to model the internal structure of stars.
- Overview of how basic stellar properties are observationally measured.
- Study the microphysics relevant for stars: the equation of state, the opacity, nuclear reactions.
- Examine the properties of simple models for stars and consider how real models are computed.
- Survey (mostly qualitatively) how stars evolve, and the endpoints of stellar evolution.

Stars are relatively simple physical systems

Problem of Stellar Structure

We want to determine the structure (density, temperature, energy output, pressure as a function of radius) of an isolated mass M of gas with a given composition (e.g., H, He, etc.)

Simplifying assumptions

- 1. No rotation \rightarrow spherical symmetry For sun: rotation period at surface \sim 1 month orbital period at surface \sim few hours
- 2. No magnetic fields

For sun: magnetic field \sim 5G, \sim 1KG in sunspots equipartition field \sim 100 MG Some neutron stars have a large fraction of their energy in B fields

3. Static

For sun: convection, but no large scale variability Not valid for forming stars, pulsating stars and dying stars.

4. Newtonian gravity

For sun: escape velocity \sim 600 km/s $<< c$ Not true for neutron stars

The Variables of Stellar Structure

Eulerian vs. Lagrangian

The Variables of Stellar Structure

Values for the Sun

$$
Mass \equiv 1M_{\odot} \approx 2 \times 10^{33} g
$$

Radius
$$
\equiv 1R_{\odot} \approx 7 \times 10^{10} \text{cm}
$$

Luminosity
$$
\equiv 1L_{\odot} \approx 4 \times 10^{33} \text{erg/s}
$$

Surface Temperature $\approx 5,800 \text{ K}$

Composition $\approx 70\%$ H 28% He 2% metals

mass in shell = density \times volume of shell

$$
dm = \rho \times 4\pi r^2 dr
$$

1. Mass Conservation

dm dr $= 4\pi r^2$ ρ

dr dm = 1 $4\pi r^2$ ρ

eulerian lagrangian

$$
P(r)4\pi r^2 - P(r+dr)4\pi r^2 - \frac{Gm}{r^2}dm = \ddot{r}dm
$$

$$
\rightarrow -dP4\pi r^2 - \frac{Gm}{r^2}dm = \ddot{r}dm \quad \rightarrow -\frac{dP}{dm}4\pi r^2 - \frac{Gm}{r^2} = \ddot{r} \quad = 0
$$

2. Hydrostatic Equilibrium

Need equation of state: $P = f(\rho, T, X_i)$

e.g., for an ideal gas:
$$
P = \frac{R}{\mu} \rho T
$$

 ε = energy generation rate per unit mass (erg/s/g)

3. Energy Generation

dLr dr $= 4\pi r^2$ ρε

dLm dm $=$ ε

eulerian lagrangian

Need nuclear physics: $\varepsilon = f(\rho, T, X_i)$

e.g., for the proton-proton chain: $\varepsilon \approx \varepsilon_0 \rho T^4$

Energy flow: Depends on opacity K (area/mass)

- radiation *low opacity*
- convection *high opacity*
- conduction *unimportant*

For radiation:

radiation pressure decreases outward \rightarrow photons have net movement outward in their random walk.

Random walk of photon through the sun.

Straight path: 2.3 seconds Random walk: 30,000 years

4. Energy Flow (radiation)

$$
\frac{dT}{dr} = -\frac{3\kappa\rho L_r}{16\pi a cr^2 T^3} \left[\frac{dT}{dm} = -\frac{3\kappa L_m}{64\pi^2 a cr^4 T^3} \right]
$$

eulerian lagrangian

Need opacity:
$$
\kappa = f(\rho, T, X_i)
$$

e.g., for electron Thomson scattering: $K \approx K_0$

4. Energy Flow (convection)

eulerian lagrangian

Need adiabatic index:
$$
\Gamma_2 = f(\rho, T, X_i)
$$

e.g., for a simple ideal gas: $\Gamma_{2} = 5/3$

Solving the Equations

Solve in 1 dimension from center to surface

Boundary Conditions

Center
$$
(r = 0)
$$
: $m = 0$, $L_r = 0$, $P = P_c$, $T = T_c$
Surface $(r = R)$: $m = M$, $L_r = L$, $P \sim 0$, $T \sim 0$

Lagrangian

Center
$$
(m = 0)
$$
: $r = 0$, $L_r = 0$, $P = ?$, $T = ?$
Surface $(m = M)$: $r = ?$, $L_r = ?$, $P \sim 0$, $T \sim 0$

Boundary conditions are incomplete at each end

Complications

1. Stars are luminous

radiate away energy \rightarrow must change in time chemical composition is changing. Must account for dX $_{\sf i}$ /dt

- 2. Convection is very complicated and important
- 3. Convection can change chemical composition
- 4. Opacities are hard to calculate. And they matter!

Homology Relations

To calculate the luminosity or radius of a star of mass *M*, we must solve these differential equations. However, we can get approximate scaling relations by using *Homology.*

Assume that each differential or local quantity simply scales with the global value of that quantity.

Homology Relations

If we also know how pressure, opacity, and nuclear generation rate scale with density and temperature, we can solve all these equations to get scaling relations. Then we can turn these scaling relations into actual equations by normalizing to the Sun.

For example, suppose we find that $L \sim M^{\alpha}$

This means that $L = const \times M^{\alpha}$

For the Sun, this is $\;\; L_{_\odot}= \mathrm{const} \times {M}_{_\odot}^{\alpha}$

Dividing the two equations we get

$$
\frac{L}{L_{\odot}} = \left(\frac{M}{M_{\odot}}\right)^{\alpha}
$$

Observations of stars

Hyades open cluster

PRC99-26 • Space Telescope Science Institute • Hubble Heritage Team (AURA/STScl/NASA)

Astronomers count photons!

We can measure three things about a photon:

- What direction did it come from?
- When did it arrive?
- What was its energy (wavelength/color)?
1. Count photons in a fixed region of the sky (aperture), a fixed window of time (exposure time), and a fixed wavelength range (filter or band).

what we call "*photometry*"

2. Count photons as a function of direction in a fixed window of time (exposure time), and a fixed wavelength range (filter or band).

what we call "*imaging*"

3. Count photons as a function of wavelength in a fixed region of the sky (aperture), and a fixed window of time (exposure time).

what we call "*spectroscopy*"

4. Count photons as a function of time in a fixed region of the sky (aperture), and a fixed wavelength range (filter or band).

what we call "*time series photometry*"

5. Count photons as a function of direction AND time in a fixed wavelength range (filter or band).

what we call "*time series imaging*"

6. Count photons as a function of wavelength AND time in a fixed region of the sky (aperture).

what we call "*time series spectroscopy*"

7. Count photons as a function of direction AND wavelength in a fixed window of time (exposure time).

what we call "*integral field spectroscopy*"

8. Count photons as a function of direction AND time AND wavelength.

the holy grail of observational astronomy…

Luminosity and flux

Luminosity L : energy/time (erg/s) Flux f : luminosity/area (erg/s/cm²)

The Electromagnetic Spectrum

SDSS filters

Photometric Filters

Astronomical fluxes are usually measured using filters

The flux in the SDSS *g* filter is:

$$
f_g = \int_0^\infty F(\lambda) S_g(\lambda) d\lambda
$$

The flux in the SDSS *r* filter is:

$$
f_r = \int\limits_0^\infty F(\lambda) S_r(\lambda) d\lambda
$$

Apparent magnitude

$$
m = -2.5 \log f + const
$$

A star that is 5 magnitudes brighter (smaller m) has 100x the flux.

$$
m_1 - m_2 = -2.5 \log(f_1/f_2)
$$

$$
\frac{f_1}{f_2} = 10^{(m_2 - m_1)/2.5}
$$

Absolute magnitude

M = apparent magnitude the star would have if it were 10pc away.

$$
f = \frac{L}{4\pi d^2} \qquad f_{10} = \frac{L}{4\pi (10 \, pc)^2}
$$

$$
m - M = -2.5 \log \left[\frac{L/4 \pi d^2}{L/4 \pi (10 pc)^2} \right] = -2.5 \log \left[\left(\frac{d}{10 pc} \right)^{-2} \right]
$$

$$
m - M = 5 \log \left(\frac{d}{10 \, pc}\right)
$$

distance modulus

Hipparcos Data

Table 3.6.4. The 150 most luminous stars in the Hipparcos Catalogue.

Bolometric magnitudes

M is measured in a band. To get the light from all wavelengths, we must add a correction.

Bolometric correction:

$$
M_{bol} = M_V + BC
$$

$$
m_{bol} = m_V + BC
$$

BC depends on band and star spectrum. By definition, BC=0 for V-band and T=6600K

Color

Color = crude, low resolution, estimate of spectral shape

$$
B - V = m_B - m_V = M_B - M_V = -2.5 \log \left(\frac{f_B}{f_V} \right)
$$

- distance independent
- indicator of surface temperature
- by definition, *(B-V)*=0 for Vega (T~9500K)

Color

• Measure a star's brightness through two different filters

• Take the ratio of brightness: (redder filter)/(bluer filter) if ratio is large \rightarrow red star if ratio is small \rightarrow blue star e.g., V/B

Color

wavelength (nm)

The color of a star measured like this tells us its temperature!

Stellar spectra

The solar spectrum can be approximated as

• a blackbody

 $+$

• absorption lines (looking at hotter layers through cooler outer layers)

Blackbody radiation

$$
B_{\nu}(T) = \frac{2h\nu^3}{c^2} \frac{1}{e^{h\nu/kT} - 1}
$$

erg s^{-1} cm $^{-2}$ Hz $^{-1}$ st $^{-1}$

 $h v_{\rm max} \sim 2.8 kT$

 \cdot Effective temperature of a star = T of a blackbody that gives the same Luminosity per unit surface area of the star.

$$
L = 4\pi R^2 \sigma T_e^4
$$
 Stefan-Boltzmann law

For sun: T_e = 5,778 K

Blackbody Spectrum or thermal spectrum

Stellar spectra are not perfect blackbodies

Atomic energy levels

- electrons orbit the nucleus in specific energy levels
- electrons can jump between energy levels given the right energy

Emission of light

Absorption of light

Energy levels for Hydrogen

Visible spectrum shows signature of hydrogen atoms

$$
E = hv = \frac{hc}{\lambda}
$$

Emission line spectrum

Absorption line spectrum

Spectrum of Sun

Spectral lines

Strength of lines depends on temperature.

e.g., Balmer lines: transitions from n=2 to higher states

Stellar spectra

Spectral lines

Lines depend on temperature in stellar atmosphere $+$ ionization potentials for relevant species e.g., H HeI HeII CaI CaII FeI 13.6eV 24.6eV 54.5eV 6.1eV 11.9eV 7.9eV Ionization occurs when $kT \sim$ ionization potential/10

Spectral classification

Spectral classification

Spectral classification

Oh Be A Fine Girl/Guy Kiss Me

Omnivorous Butchers Always Find Good Kangaroo Meat

Only Bored Astronomers Find Gratification Knowing Mnemonics
Luminosity class

Stars of same type have different line widths

Same *T*, different $R \longrightarrow$ different surface gravity $g \longrightarrow$ **→ different surface pressure P**

Pressure broadening: orbitals of atoms are perturbed due to collisions \longrightarrow broadening of spectral lines.

 \mathbf{S} ince $L = 4\pi R^2 \sigma_{_T} T^4$, changes in R at fixed T are changes in L

Spectral line widths \longrightarrow luminosity classification

Spectra of AO stars $3,5$ AO V $-$ AO III $\overline{3}$ $AO I$ $2,5$ $\overline{2}$ $1,5$ $\,1\,$ 0.5 \triangle 4000 5000 6000 7000 8000 9000 10000 Wavelength (Angstroms)

Special stars

- C: carbon stars same T_{eff} as K, M stars, but higher abundance of C than $O \rightarrow$ all O goes to form CO. Remaining C forms C2, CN.
- S: same T_{eff} as K, M stars, but have extra heavy elements
- W: Wolf-Rayet He in atmosphere instead of H, strong winds
- L: cooler than M stars. Some do not have fusion.
- T: cool brown dwarfs (700-1,000K). Methane lines are prominent.

The Sun is a G2V star

The first Hertzsprung-Russell (H-R) diagram

Figure 8.10 Henry Norris Russell's first diagram, with spectral types listed along the top and absolute magnitudes on the left-hand side. (Figure from Russell, Nature, 93, 252, 1914.)

Hipparcos Color-Magnitude Diagram

Plot luminosity vs. temperature

 $L = 4\pi R^2 \sigma T^4$

Hipparcos H-R diagram

Theoretical H-R diagram

Chemical composition

Primordial (Big Bang) nucleosynthesis: protons fuse to form He and heavier elements 3 minutes after the Big Bang. Ends 20 minutes later. Alpher, Bethe & Gammow, Physical Review L, 1948

75% H 25% He 0.01% D

Subsequent fusion inside massive stars and enrichment of the inter-stellar medium via supernovae, leads to future generations of stars with more heavy elements.

Stellar populations in the Milky Way

Stellar populations in the Milky Way

Determining stellar properties

Determining distance

July Earth Star B *R* R_1^{\dagger} $\tan \pi =$ $\approx \pi$ $\left(\begin{array}{c} \Lambda_1 \\ \vdots \\ \Lambda_m \end{array}\right)$ π *d d* Star A **A.U.** 149.6 10 Star C $R = 1AU = 1.5 \times 10^{13}$ cm January

Define new distance unit: parsec (**par**allax-**sec**ond)

$$
1pc = \frac{1AU}{\tan(1'')} = 206,265AU = 3.26ly
$$

RULER

Point spread function (PSF)

Need high angular precision to probe far away stars.

$$
d=\frac{1}{\pi}
$$

Error propagation:

$$
\sigma_d = \sqrt{\left(\frac{\partial d}{\partial \pi}\right)^2 \sigma_\pi^2} = \sqrt{\left(-\frac{1}{\pi^2}\right)^2 \sigma_\pi^2} = \frac{\sigma_\pi}{\pi^2} = \frac{\sigma_\pi}{\pi} d
$$

$$
\frac{\sigma_d}{d} = \frac{\sigma_\pi}{\pi}
$$

At what distance do we get a given fractional distance error?

$$
d = \left(\frac{\sigma_d}{d}\right) \frac{1}{\sigma_{\pi}}
$$

e.g., to get 10% distance errors and

$$
d_{\max} = \frac{0.1}{\sigma_{\pi}}
$$

Determining distance: moving cluster method

$$
\left(\frac{d}{1pc}\right) = \frac{\left(v_t/1 \text{ kms}^{-1}\right)}{4.74 \left(\mu/1'' \text{yr}^{-1}\right)}
$$

 $d =$

 v_t

 μ

Determining distance: moving cluster method

- 1. Measure proper motions of stars in a cluster
- 2. Obtain convergent point
- 3. Measure angle between cluster and convergent point θ
- 4. Measure radial velocity of cluster V_r
- 5. Compute tangential velocity V_t
- 6. Use proper motion of cluster to get distance *d*

$$
d = \frac{v_r \tan \theta}{4.74 \,\mu}
$$

RULER

Determining distance: secular parallax

- Parallax method is limited by 2AU baseline of earth's orbit
- Sun moves ~4AU/yr toward Vega relative to local rotation of Galactic disk
- Over a few years, this can build up to a large baseline
- Unfortunately, other stars are not at rest, rather have unknown motions
- However, if we average over many stars, their mean motion should be zero (relative to local rotation)
- Can therefore get the mean distance to a set of stars

Determining distance: secular parallax

RULER

Determining distance: Baade-Wesselink (moving stellar atmosphere)

CANDLE

 R_{2}

For pulsating stars, SN, novae, measure flux and effective temperature at two epochs, as well as the radial velocity and time between the epochs.

$$
f_1, f_2, T_1, T_2, v_r(t), \Delta t
$$

$$
\frac{f_1}{f_2} = \frac{L_1}{L_2} = \left(\frac{R_1}{R_2}\right)^2 \left(\frac{T_1}{T_2}\right)^4 \longrightarrow \frac{R_1}{R_2} = \left(\frac{f_1}{f_2}\right)^{\frac{1}{2}} \left(\frac{T_2}{T_1}\right)^2
$$

$$
R_2 - R_1 = \overline{v}_r \times \Delta t
$$

Solve for R₁ and R₂. R and T $\longrightarrow L \longrightarrow d$

Determining distance: spectroscopic parallax

- Stellar spectra alone can give us the luminosity class + spectral class \rightarrow position on HR diagram
- This gives the absolute magnitude, which gives the distance modulus.
- Basically, compare a star's spectrum to an identical spectrum of another star with known luminosity.
- This method is poor in accuracy (+/- 1 magnitude error in absolute magnitude \rightarrow 50% error in distance)
- But it can be applied to all stars

Determining distance: main sequence fitting

- Measure the colors and magnitudes of stars in a cluster (e.g., *r* and *g-r*)
- Plot the HR diagram: *r* vs. *g-r*
- Compare the the HR diagram of another cluster of known distance: *Mr* vs. *g-r*
- Find the vertical offset in HR diagram between the main sequences of the two clusters \rightarrow distance modulus

$$
m - M = 5\log d - 5
$$

Determining distance: main sequence fitting

Determining distance: variable stars

Cepheid variables:

Pop I giants, $M \sim 5{\text -}20 \text{ M}_{\text{sun}}$

Pulsation due to feedback loop:

An increase in T

- \rightarrow HeIII (doubly ionized He)
- \rightarrow high opacity
- \rightarrow radiation can't escape
- \rightarrow even higher T and P
- \rightarrow atmosphere expands
- \rightarrow low T
- \rightarrow HeII (singly ionized He)
- \rightarrow low opacity
- \rightarrow atmosphere contracts
- \rightarrow rinse and repeat...

Data from a Well-Measured Cepheid

Time (usually Days)

Determining distance: variable stars

RR-Lyrae variables: Pop II dwarfs, $M \sim 0.5 M_{sun}$

Apparent V magnitude of variable star RR Lyr

Determining distance: variable stars

Variable stars have a tight period-luminosity relation

- Measure lightcurves: flux(t)
- Get period P
- From P-L relation, get L
- Use L to get distance

Very powerful method. Cepheids can be seen very far away. Used to measure H_0

P-L relation is calibrated on local variables with parallax measurements

PERIOD - LUMINOSITY RELATIONSHIP

Determining distance: dynamical parallax

For binary star systems on main sequence

- Measure: period of orbit P, angular separation $\,\theta$, fluxes f_1 and f_2
- Kepler's 3rd law: $P^2 =$ $4\pi^2$ $G(M_1 + M_2)$ *a* 3

• Assume
$$
M_1 = M_2 = M_{\odot}
$$

- Use Kepler to get *a*
- Use θ and a to get preliminary distance
- Use distance and fluxes to get luminosities L_1 , L_2
- Use mass-luminosity relation for main sequence to get better masses
- Iterate until convergence

Determining Luminosity

1. Measure flux or magnitude Measure distance $L = 4\pi d^2 f$ $M = m - 5 \log d - 5$

T, *L*

2. Measure spectrum

line ratios and widths (i.e., compare the star to a star of identical spectrum that has a known L)

Determining Temperature

- 1. Spectral lines \rightarrow spectral type
- 2. Colors (cheap and accurate)
- 3. Blackbody fitting or Wien's law: *h*^ν ∼ 2.8*kT*(not good for very hot or cool stars)
- 4. Stellar atmosphere modeling (uncertain)
- 5. Measure angular size and flux (need very high resolution imaging)

 $R = \theta_R d$ $L = 4\pi d^2 f$

$$
L = 4\pi R^2 \sigma T^4 \rightarrow T = \left(\frac{L}{4\pi R^2 \sigma}\right)^{\frac{1}{4}} \rightarrow T = \left(\frac{4\pi d^2 f}{4\pi \left(\theta_R d\right)^2 \sigma}\right)^{\frac{1}{4}}
$$

$$
T = \left(\frac{f}{\sigma \theta_R^2}\right)^{\frac{1}{4}}
$$

Determining Radius

- Very difficult because angular sizes are tiny. The sun at 1pc distance has an angular radius of ~0.5mas
- Important because of:

• surface gravity
$$
g \sim \frac{GM}{R^2}
$$

• density
$$
\rho \sim \frac{M}{R^3}
$$

• Temperature
$$
T \sim (L/R^2)^{1/4}
$$

• testing models

Determining Radius: Interferometry

Interferometry yields high resolution images

1. Speckle interferometry

 ~ 0.02 as

Diffraction limited:

$$
\theta = \frac{1.22 \lambda}{D_T}
$$

e.g., for HST (D_T=2.4m), θ =0.05 as

Determining Radius: Interferometry

 \sim 0.5 mas

The sun's radius could be measured out to \sim 1-10pc

Determining Radius

- Luminosity + Temperature \rightarrow Radius
- Baade-Wesselink (moving atmosphere)
- Lunar Occultation

As a star disappears behind the moon, a Fresnel diffraction pattern is created that depends slightly on $\,\theta_{\scriptscriptstyle R}^{}$

 \sim 2 mas, only good for stars in ecliptic.

Determining radius: eclipsing binaries

Determining radius: eclipsing binaries

Determining Mass

Only possible to measure accurately for binaries.

Kepler's 3rd law: $G(M_1 + M_2)P^2 = 4\pi^2 a^3$

 \blacktriangledown

$$
M_1 + M_2 \approx M_0
$$

For Sun-Earth system: $P = 1yr$ \longrightarrow $GM_{\odot} (1yr)^2 = 4\pi^2 (1AU)^3$
a=1AU

$$
\left[\left(\frac{M_1 + M_2}{M_\odot} \right) \left(\frac{P}{1 \text{yr}} \right)^2 = \left(\frac{a}{1 \text{AU}} \right)^3 \right]
$$

Determining Mass: visual binaries

Visual binaries are binaries where the angular separation is detectable and the orbit can be traced out.

• Find the center of mass of the system (c.o.m. must move with constant velocity)

$$
M_1 a_1 = M_2 a_2 \rightarrow \frac{M_1}{M_2} = \frac{a_2}{a_1} = \frac{\alpha_2}{\alpha_1}
$$

• Measure the period, semi-major angular separation and distance.

$$
\begin{array}{ccc}\n\alpha = \alpha_1 + \alpha_2 & & \\
a = d\alpha & & \\
M_1 + M_2 = \frac{(d\alpha)^3}{P^2}\n\end{array}
$$

- Solve for individual masses. Very sensitive to distance errors: $M \sim d^3$
- If distance is not known, radial velocity data is sufficient to get $|a|$ e.g., for a circular orbit: $v = 2\pi a / P$

Determining Mass: visual binaries

If the orbit is inclined, then we must know the inclination angle *i*

$$
\frac{M_1}{M_2} = \frac{a_2}{a_1} = \frac{\alpha_2 \cos i}{\alpha_1 \cos i} = \frac{\alpha_2}{\alpha_1}
$$
 Don't need *i*

$$
M_1 + M_2 = \left(\frac{\alpha}{\cos i}\right)^3 \frac{d^3}{P^2}
$$
 Need *i*

In an inclined orbit, the center of mass does not lie at the ellipse focus. Find the projection that fixes this $\rightarrow \cos i$

Spectroscopic binaries are unresolved binaries that are only detected via doppler shifts in their spectrum.

- Single line binaries: single set of shifting spectral lines
- Double line binaries: two sets of spectral lines (one from each star)
- Two sets of lines, but for different spectral types

Determining mass: spectroscopic binaries

Determining Mass: spectroscopic binaries

• Can get a lower limit on M₁+M₂ since sin *i* <1

Determining Mass: spectroscopic binaries

Need to measure both radial velocities - only possible with double line binaries.

Need to know sin *i*. This is possible when:

- also an eclipsing binary: sin *i* = 1
- also a visual binary: measure orbit
- compute M₁+M₂ for a statistical sample where <sin³*i*> is known (For an isotropic distribution, <sin3*i*>=0.42. However, no Doppler shift will be observed if *i*=0, so there is a selection effect that favors high inclinations)

Determining Mass: surface gravity

- Pressure broadening of spectral lines \rightarrow surface gravity
- $g =$ *GM* R^2

• Measure radius $R \rightarrow$

$$
M=\frac{gR^2}{G}
$$

• Sensitive to errors in R

Determining Mass: Gravitational lensing

Discovery of Widest-separation Quasar
Gravitational Lens

Suprime-Cam (g',r',i')
December 17, 2003

Subaru Telescope, National Astronomical Observatory of Japan
Copyright © 2003, National Astronomical Observatory of Japan, All rights reserved.

Determining Mass: Gravitational lensing

Determining Mass: gravitational microlensing

Case of lensing when multiple images are unresolved and we only detect an increase in the flux of one image.

Determining mass: gravitational microlensing

Determining mass: gravitational microlensing

Determining mass: gravitational microlensing

Determining Chemical Composition

Theoretical: fractional abundances by mass

X: Hydrogen Y: Helium Z: metals Sun: (0.7, 0.28, 0.02)

Observational: number density relative to Hydrogen, normalized to sun.

e.g., iron abundance
$$
[Fe/H] = \log \left(\frac{n(Fe)}{n(H)} \right)_{*} - \log \left(\frac{n(Fe)}{n(H)} \right)_{\odot}
$$

So, for solar abundance: $\left [\mathrm{Fe}/\mathrm{H} \right] = 0$, 10 times more iron: $\left [\mathrm{Fe}/\mathrm{H} \right] = +1$ In our galaxy, $\rm [Fe/H]$ ranges from -4.5 to +1.

Need high resolution spectra + stellar atmosphere models to get abundances via fitting.

Hipparcos stars Determining age: Main Sequence turn-off

The lack of blue dwarfs tells us that the age of M6 is sufficiently old that the massive blue stars have died.

Yellow and red stars are still present because they live longer lives.

Determining age: Main Sequence turn-off

- All stars arrived on the mainsequence (MS) at about the same time.
- The massive stars on the top left of MS are the first to go.
- The cluster is as old as the most luminous star that remains on the MS.
- The position of the hottest, brightest star on a cluster's MS is called the *main-sequence turnoff point*.

Determining age: Main Sequence turn-off

Which cluster is oldest?

NGC 188: 7 billion years

 $B-V$

 $V-R$

Measurement Errors

- errors in brightness/colors
- errors in distance/luminosity
- errors in cluster membership

Theoretical Errors

- uncertain chemical abundances
- uncertain amount of dust
- uncertain stellar physics

Typical error < 1 billion years

Global Energetics

Total stellar energy

 $W =$ gravitational potential energy Ω + kinetic energy due to bulk motions (turbulence, pulsation) + internal energy from microscopic processes *U*
Gravitational Potential Energy

Gravitational potential energy

 Ω : energy required to assemble the star by collecting material from the outside universe.

= negative of energy required to disperse the star.

$$
d\Omega = -\int_{r}^{\infty} \frac{Gmdm}{r'^2} dr' = -\frac{Gm}{r} dm
$$

$$
\Omega = -\int_{0}^{M} \frac{Gm}{r} dm
$$

which depends on the density profile $\rho(r)$

Gravitational Potential Energy

e.g., for
$$
\rho(r) = \rho_0
$$
 $\begin{array}{c} m = \rho_0 \frac{4}{3} \pi r^3 \\ M = \rho_0 \frac{4}{3} \pi R^3 \end{array}$ $\rightarrow \frac{m}{M} = \left(\frac{r}{R}\right)^3 \rightarrow r = R \left(\frac{m}{M}\right)^{\frac{1}{3}}$

$$
\Omega = -\int_{0}^{M} \frac{Gm}{r} dm = -\frac{GM^{1/3}}{R} \int_{0}^{M} m^{2/3} dm = -\frac{3}{5} \frac{GM^{2}}{R}
$$

In general,
$$
\Omega = -q \frac{GM^2}{R}
$$
 and $q > \frac{3}{5}$ since $\rho \downarrow$ with r

$$
\rho \sim r^{-1} \to q = \frac{2}{3} \qquad \rho \sim r^{-2} \to q = 1
$$

Internal and Kinetic Energy

Internal energy

U : Kinetic energy due to motions of particles (thermal, not bulk) + energy in atomic and molecular bonds.

$$
U = -\int_{M} E_{m} dm
$$
 Where E_{m} : local specific internal energy (erg/g)

Kinetic energy

 $K:$ Total kinetic energy (thermal + bulk)

Consider all particles in a star at positions \vec{r}_i with momenta $\overrightarrow{1}$ *ri* \Rightarrow \vec{p}_i

$$
Q = \sum_{i} \vec{p}_i \cdot \vec{r}_i
$$

evaluate $\frac{dQ}{dt}$
what are the units of $\frac{dQ}{dt}$? (energy units: $\frac{m \cdot v \cdot r}{t} = m \cdot v^2$)

t

 $Q = \sum \vec{p}_i$. \rightarrow $\sum \vec{p}_i \cdot \vec{r}_i$ *i*

$$
\frac{dQ}{dt} = \frac{d}{dt} \sum_{i} m \cdot \dot{\vec{r}}_{i} \cdot \vec{r}_{i}
$$
\n
$$
= \frac{d}{dt} \sum_{i} \frac{d}{dt} \left(\frac{1}{2} m \cdot \vec{r}_{i}^{2} \right)
$$
\n
$$
= \frac{1}{2} \sum_{i} \frac{d^{2}}{dt^{2}} \left(m \cdot \vec{r}_{i}^{2} \right)
$$

m ⋅ \Rightarrow $\sum m \cdot \vec{r}_i^2$ *i*

is just the total moment of inertia of the system.

dQ dt = 1 2 d^2I dt^2

 $Q = \sum \vec{p}_i$. \rightarrow *ri i* ∑

$$
\frac{dQ}{dt} = \sum_{i} \frac{d\vec{p}_i}{dt} \cdot \vec{r}_i + \sum_{i} \vec{p}_i \cdot \frac{d\vec{r}_i}{dt}
$$

x total kinetic energy)

$$
\begin{aligned} \boxed{\mathbf{B} :} \quad \sum_{i} \vec{p}_{i} \cdot \frac{d\vec{r}_{i}}{dt} \; &= \; \sum_{i} m \cdot \vec{v}_{i} \cdot \vec{v}_{i} \\ &= \; 2 \sum_{i} \frac{1}{2} m \cdot \vec{v}_{i}^{2} \; \underbrace{\vert \; = \; 2K \vert}_{\; \;} \end{aligned} \tag{2}
$$

$$
\begin{aligned}\n\boxed{\mathbf{A:}} \quad & \sum_{i} \frac{d\vec{p}_{i}}{dt} \cdot \vec{r}_{i} \ = \ \sum_{i} \frac{d}{dt} \Big(m \cdot \dot{\vec{r}}_{i} \Big) \cdot \vec{r} \quad = \ \sum_{i} m \cdot \ddot{\vec{r}}_{i} \cdot \vec{r}_{i} \quad = \ \sum_{i} \vec{F}_{i} \cdot \vec{r}_{i} \\
\vec{F}_{i} \quad \text{is the sum of all gravitational forces on particle } i \quad = \ \sum_{j \neq i} \vec{F}_{ij} \\
\boxed{\vec{r}_{i} \\
\boxed{\vec{r}_{j} - \vec{r}_{i} \\
\boxed{\vec{r}_{j} - \vec{r}_{i}}} \quad \vec{F}_{ij} = G \frac{m_{i} m_{j}}{r_{ij}^{2}} \frac{\Big(\vec{r}_{j} - \vec{r}_{i} \Big)}{r_{ij}} \ = \ -G \frac{m_{i} m_{j}}{r_{ij}^{3}} \Big(\vec{r}_{i} - \vec{r}_{j} \Big) \\
\boxed{\vec{r}_{j} \\
$$

energy r_{ij}^3 r_{ij} $i \quad j > i$

j>*i i*

$$
\frac{1}{2}\frac{d^2I}{dt^2} = 2K + \Omega
$$

In a globally static system:

$$
\frac{d^2I}{dt^2} = 0 \quad \longrightarrow \quad \boxed{2K = -\Omega}
$$

For a collisionless system:
$$
W = K + \Omega
$$
 \longrightarrow $W = \frac{1}{2}\Omega = -K$

bound!

What about a collisional system? How does the 2*K* term relate to the internal energy of a gas?

Kinetic theory: pressure in a gas is caused by the sum of collisions of particles and momentum transfer

Particle *i* colliding with wall in x-direction

$$
\Delta p_i = 2 p_{x,i} = 2 m_i v_{x,i}
$$

Time between collisions of same particle with wall

$$
\Delta t = \frac{2l}{v_{x,i}}
$$

Force due to particle i

$$
F_i = \frac{\Delta p_i}{\Delta t} = \frac{m_i v_{x,i}^2}{l}
$$

Total force on wall

$$
F = \sum_{i} \frac{m_{i}v_{x,i}^{2}}{l} = \frac{1}{l} \sum_{i} m_{i}v_{x,i}^{2}
$$

Kinetic theory: pressure in a gas is caused by the sum of collisions of particles and momentum transfer

i

$$
P \text{ressure}
$$
\n
$$
P = \frac{F}{A} = \frac{2K}{3l \cdot A} = \frac{2K}{3V} \longrightarrow \boxed{2K = 3P \cdot V}
$$

 $m_i v_i^2$

i

In reality, pressure is changing over volume (can think of our box as a volume element *dV*)

$$
2K = 3\int\limits_V P dV
$$

In a star's spherical shells

$$
dm = \rho dV \qquad 2K = \int_{M} \frac{3P}{\rho} dm
$$

Depends on the equation of state

For a relation btw *P* and E_m of $\ P=(\gamma-1)\rho E_m-\gamma$ -law equation of state)

$$
2K = \int_{M} 3(\gamma - 1) E_m \, dm = 3(\gamma - 1) \int_{M} E_m \, dm = 3(\gamma - 1) U
$$

K is only equal to U if $\gamma = \frac{3}{2}$ — i.e., for an ideal monoatomic gas 5 3

Virial theorem:

$$
3(\gamma - 1)U + \Omega = 0
$$

Total energy

 W

$$
= \Omega + U
$$

=
$$
\Omega - \frac{\Omega}{3(\gamma - 1)}
$$

=
$$
\Omega \left(1 - \frac{1}{3(\gamma - 1)}\right)
$$

$$
=\Omega \frac{3\gamma-4}{3(\gamma-1)}
$$

Total energy must be ≤ 0 for star to be stable. Otherwise, it has enough energy to disperse itself.

$$
\Rightarrow \gamma > \frac{4}{3}
$$

• Ideal gas $\left(\gamma = \frac{5}{3}\right)$: safely bound
• Radiation $\left(\gamma = \frac{4}{3}\right)$: unstable

As radiation pressure becomes more important, star becomes less bound.

As a star radiates, it loses energy (*without other energy sources*).

$$
\dot{W} < 0 \rightarrow \begin{cases} \dot{\Omega} < 0 \\ \dot{U} > 0 \end{cases} \rightarrow \text{star contracts}
$$

and heats up

The Virial Theorem: Applications

5

Internal temperature

- For an ideal monoatomic gas:
- Also, the energy density is:

$$
\gamma = \frac{5}{3} \rightarrow U = -\frac{\Omega}{2}
$$

$$
E = \frac{3}{2}nkT \rightarrow U = \frac{3}{2}nkTV
$$

 $3(\gamma - 1)U + \Omega = 0$

n : number density of particles μ $N_{\overline{A}}$: : \vert ⎨ \int \lfloor ⎪ mean molecular weight Avogadro's number $\begin{array}{c} \hline \end{array}$ $\left\{ \right.$ $\overline{\mathcal{L}}$ \int ⎪ $n =$

$$
n=\frac{\rho N_A}{\mu}
$$

$$
U = \frac{3}{2} \frac{\rho N_A}{\mu} kTV = \frac{3}{2} \frac{N_A}{\mu} kMT \rightarrow T = \frac{2}{3} \frac{\mu}{N_A k} \frac{U}{M}
$$

• Virial theorem: $U = -\frac{\Omega}{2} = \frac{q}{2} \frac{GM^2}{R}$

The Virial Theorem: Applications

Internal temperature

$$
T = \frac{q}{3} \frac{\mu G}{N_A k} \frac{M}{R}
$$

$$
T = 5 \times 10^6 K \left(\frac{q}{3/5}\right) \left(\frac{M}{M_{\odot}}\right) \left(\frac{R}{R_{\odot}}\right)^{-1}
$$

(actual central temperature of sun is 15 million K)

- Internal $T \gg$ surface $T \rightarrow$ strong T gradient
- Most of the sun is highly ionized
- T is sufficiently high for nuclear fusion (~1 million K)
- Fusion happens due to gravitational energy

Dynamical timescale

• Virial theorem:
$$
\frac{d^2I}{dt^2} \approx \frac{GM^2}{R}
$$

$$
\frac{1}{2}\frac{d^2I}{dt^2} = 2K + \Omega
$$

$$
\frac{I}{t_{\text{dyn}}^2} \approx \frac{GM^2}{R} \rightarrow t_{\text{dyn}} \approx \left(\frac{R \cdot I}{GM^2}\right)^{1/2} \rightarrow t_{\text{dyn}} \approx \left(\frac{R^3}{GM}\right)^{1/2} \approx (G\rho)^{-1/2}
$$

$$
I \approx MR^2
$$

*v*esc ² *GM R ^t*dyn *^R v*esc • Escape velocity: • Timescale for collapse: *^R GM R* ⎛ ⎝ [⎜] [⎞] ⎠ ⎟ ¹ ² *t*dyn (*G*ρ) −1 2 For sun, *t*dyn~ 1 hour

Kelvin-Helmholtz timescale

Timescale for star to radiate away its thermal/gravitational energy.

• Star contracts slowly maintaining hydrostatic equilibrium

$$
\Delta R < 0
$$
\n
$$
\Omega = -q \frac{GM^2}{R}
$$
\n
$$
\Delta \Omega < 0
$$
\n
$$
\frac{d\Omega}{dR} = q \frac{GM^2}{R^2}
$$
\n
$$
\Delta \Omega = +q \frac{GM^2}{R^2} \Delta R
$$

• Virial theorem:
$$
\gamma = \frac{5}{3} \rightarrow W = \frac{\Omega}{2} \rightarrow \Delta W < 0
$$

• Energy is lost from the system: radiation

 $\Delta W =$ $ΔΩ$ half goes to increasing *U* and half goes to luminosity

Kelvin-Helmholtz timescale Timescale for star to radiate away its thermal/gravitational energy.

• Suppose contraction is solely responsible for maintaining luminosity

$$
L = -\frac{dW}{dt} = -\frac{d}{dt} \left(\frac{\Omega}{2}\right) = -\frac{1}{2} \frac{d\Omega}{dR} \frac{dR}{dt} = -\frac{q}{2} \frac{GM^2}{R^2} \frac{dR}{dt}
$$

$$
t_{\text{KH}} = \frac{q \, GM^2}{2 \, LR}
$$

$$
t_{\rm KH} = 2 \times 10^7 \,\text{yr} \left(\frac{M}{M_{\odot}}\right)^2 \left(\frac{L}{L_{\odot}}\right)^{-1} \left(\frac{R}{R_{\odot}}\right)^{-1}
$$

• Too short a timescale to have happened.

Nuclear timescale Timescale for star to burn its H fuel and leave the Main Sequence.

$$
t_{\text{nuc}} = \frac{\text{total available energy}}{\text{luminosity}}
$$

$$
t_{\rm nuc} = \frac{\varepsilon f_{\rm core} X M c^2}{L}
$$

 \mathcal{E} = efficiency of nuclear burning (4¹H \rightarrow ⁴He) ~ 0.007

 f_{core} = mass fraction of core (star leaves main sequence after core burns) \sim 0.1

For sun, t_{nuc} 10 billion years

For stars of solar mass or greater:

$$
\frac{L}{L_{\odot}} \approx \left(\frac{M}{M_{\odot}}\right)^{3.5}
$$

$$
t_{\rm nuc} \approx 10^{10} \,\text{yr} \left(\frac{M}{M_{\odot}}\right)^{-2.5}
$$

$$
t_{\text{dyn}} \sim 1 \text{ hour}
$$

$$
t_{\text{KH}} \sim 20 \text{ million years}
$$

$$
t_{\text{nuc}} \sim 10 \text{ billion years}
$$

$$
t_{\rm dyn} \ll t_{\rm KH} \ll t_{\rm nuc}
$$

It is valid to assume that stars on the main sequence are in dynamical and thermal equilibrium.

$$
P = f(\rho, T, X_i)
$$

To derive the E.O.S., we need to consider:

- Quantum statistics (bosons vs. fermions)
- Non-relativistic vs. relativistic particles
- Degenerate vs. non-degenerate matter

Radiation pressure and degenerate relativistic electrons

masses of stars

$$
\gamma = \frac{4}{3}
$$
 (star is not bound) This sets limits on
the masses of stan

For most normal stars, ideal gas + radiation pressure are the important contributions

$$
P = P_{\rm gas} + P_{\rm rad} \qquad = nkT + \frac{1}{3}aT^4
$$

n is the number density of all atoms and free electrons

To calculate the number density *n*, we need to know:

- constituents of plasma
- state of ionization

Available particles: ions + free electrons

$$
n = n_I + n_e = \sum_i \left(n_{I,i} + n_{e,i} \right)
$$

Each species *i* :

$$
A_i
$$
: nuclear mass number $\begin{cases} e.g., A_{i_H} = 1, A_{i_{He}} = 4, A_{i_{2_C}} = 12 \\ Z_i : \text{ nuclear charge} \\ X_i : \text{fraction by mass} \end{cases}$ usually $Z_i = \frac{A_i}{2}$ for metals

 1 mole = amount that has as many atoms of substance as there are atoms of *12C* in *12g* of *12C*.

Avogadro's number: $N_A = 6 \times 10^{23}$

e.g., 1 mole of ¹²C weighs
$$
12g
$$

1 mole of ¹H weighs $1g$
1 mole of ⁴He weighs $4g$

Ai NA # of atoms per mole = mass of mole = mass per atom (in grams)

$$
n_{I} = \frac{\rho N_{A}}{\mu_{I}}
$$

$$
\mu_I^{-1} = \sum_i \frac{X_i}{A_i}
$$

electrons

also need to know ionization fraction y_i

$$
n_{e,i} = n_{I,i} \cdot Z_i \cdot y_i = \rho N_A \frac{X_i}{A_i} Z_i y_i
$$

$$
n_e = \sum_i n_{e,i} = \rho N_A \sum_i \frac{X_i Z_i y_i}{A_i}
$$

$$
y_i = \begin{cases} 0: \text{ completely neutral} \\ 1: \text{ completely ionized} \end{cases}
$$

$$
n_e = \frac{\rho N_A}{\mu_e}
$$

Mixture of ions + electrons

$$
n = \frac{\rho N_A}{\mu}
$$

$$
\mu_e^{-1} = \sum_i \frac{X_i Z_i y_i}{A_i}
$$

 μ is the mean weight of particles in units of 1H

$$
\mu^{-1} = \mu_I^{-1} + \mu_e^{-1}
$$

$$
({}^{1}H, {}^{4}He, \text{ metals}) \equiv (X,Y,Z)
$$

 \vert

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Composition of Sun

- $^{16}O \sim 0.01$
- $^{12}C \sim 0.004$
- $56Fe \sim 0.0015$
	- $^{28}Si \sim 0.001$
	- $^{14}N \approx 0.001$
- $^{24}Mg \sim 0.0008$
- ^{20}Ne ~ 0.0006
	- $32S \sim 0.0004$

In the core

 $^{1}H \sim 0.34$ ^{4}He ~ 0.64

These were fused inside stars

 $X_i Z_i y_i$

 \overrightarrow{i} *A_i*

$$
\mu_I^{-1} = \sum \frac{X_i}{\Lambda} \bigg| \qquad \qquad \bigg| \mu_e^{-1} = \sum \frac{X_i Z_i y_i}{\Lambda} \bigg| \qquad \qquad \boxed{\mu^{-1} = \mu_I^{-1} + \mu_e^{-1}}
$$

• Fully ionized gas with zero metals $(Z=0, y_i=1)$

 $\mu_{_{e}}$

 $\frac{-1}{I}$ =

Xi

 $\overline{}$ *i* A_i

∑

 $^{-1}_{e}$ =

 $X_i Z_i y_i$

 \overrightarrow{i} *A_i*

$$
\mu_I^{-1} = \sum \frac{X_i}{A} \left| \n\mu_e^{-1} = \sum \frac{X_i Z_i y_i}{A} \right| \n\mu_I^{-1} = \mu_I^{-1} + \mu_e^{-1}
$$

• In the core, all
$$
H
$$
 is converted to He ($X=0$)

 $\mu_{_{e}}$

 $^{-1}_{e}$ =

$$
X = 0 \rightarrow \mu = 1.3
$$

• Adding metals (fully ionized)

 $\frac{-1}{I}$ =

Xi

 $\overline{}$ *i* A_i

∑

$$
\mu_{I}^{-1} = \frac{X}{A_{H}} + \frac{Y}{A_{He}} + \frac{Z}{A_{Z}} = X + \frac{Y}{4} + \frac{Z}{A_{Z}}
$$
\n
$$
\mu_{e}^{-1} = \frac{X \cdot Z_{H} \cdot y_{H}}{A_{H}} + \frac{Y \cdot Z_{He} \cdot y_{He}}{A_{He}} + \frac{Z \cdot \frac{A_{Z}}{2} \cdot y_{Z}}{A_{Z}} = X + \frac{Y}{2} + \frac{Z}{2}
$$
\n
$$
\mu^{-1} = 2X + \frac{3}{4}Y + \frac{Z}{A_{Z}} + \frac{Z}{2} = 2X + \frac{3}{4}Y + \frac{1}{2}Z
$$
\n
$$
Z = 1 \rightarrow \mu = 0.5
$$

Equations of state may be derived assuming Local Thermodynamic Equilibrium (LTE) At any position in the star, thermodynamic equilibrium holds locally even though it does not hold globally.

Particle-particle and photon-particle mean free paths are short relative to other length/time scales.

For example, the pressure scale height (height over which the pressure changed by a factor of *e*) is much longer than the mean free path.

$$
\lambda_{P} = -\left(\frac{d\ln P}{dr}\right)^{-1} = -\left(\frac{1}{P}\frac{dP}{dr}\right)^{-1} = \frac{P}{g\rho} \sim R \text{ in most of star}
$$

Compare that to the mean free path $\lambda_{\gamma} \sim 1 \text{cm}$

Distribution Functions

Statistical mechanics gives us the phase-space density of a particle species.

$$
n(p) = \frac{1}{h^3} \sum_j \frac{g_j}{\exp[(\varepsilon_j + \varepsilon(p) - \mu)/kT] \pm 1} \Delta x \Delta y \Delta z \Delta p_x \Delta p_y \Delta p_z
$$

$$
\Delta x \Delta y \Delta z \Delta p_x \Delta p_y \Delta p_z = h^3
$$

- $p = |\vec{p}|$: momentum
- g_j : degeneracy of state *j* (# of states having same energy)
- ε _{*j*} : energy of state *j* relative to some reference level
- $\varepsilon\bigl(p\bigr)$: kinetic energy $\varepsilon\bigl(p\bigr)\!=\!\left(p^2c^2+m^2c^4\right)^{\!1\!}/2$ \bullet $\bm{\varepsilon}(p)$ $\;$: kinetic energy $\;$ $\bm{\varepsilon}(p)\!=\!\left(p^2c^2+m^2c^4\right)^{\!\!1/2}-mc^2$

$$
\begin{aligned}\n\text{NR:} \left[pc \ll mc^2 \right] &= \left(1 + \frac{p^2 c^2}{m^2 c^4} \right)^{1/2} mc^2 - mc^2 \approx \left(1 + \frac{1}{2} \frac{p^2 c^2}{m^2 c^4} \right) mc^2 - mc^2 \approx \frac{p^2}{2m} \\
\text{UR:} \left[pc \gg mc^2 \right] &= \left(p^2 c^2 + m^2 c^4 \right)^{1/2} - m c^2 \approx pc \\
\text{Var:} \left[\frac{\partial E}{\partial N} \right] &:\text{chemical potential} \quad \sum \mu_i dN_i = 0 \qquad \text{e.g., } H^+ + e^- \to H^0 + \gamma\n\end{aligned}
$$

•
$$
\mu = \left(\frac{\partial E}{\partial N}\right)
$$
: chemical potential $\sum_{i} \mu_{i} dN_{i} = 0$ e.g., $H^{+} + e^{-} \rightarrow H^{0} + \gamma$
\n• \pm : + for fermions (Fermi-Dirac), - for bosons (Bose-Einstein)

Distribution Functions

We can get equation of state quantities from *n(p)* Integrate over all phase space assuming spherical symmetry

• Space density

$$
n = \int_{0}^{\infty} n(p) 4\pi p^2 dp
$$

• Internal energy (per unit volume)

$$
E = \int_{0}^{\infty} \varepsilon(p) n(p) 4\pi p^2 dp
$$

• Pressure

Kinetic theory

$$
P \cdot V = \frac{1}{3} \sum_{i} m_i v_i^2 \rightarrow P = \frac{1}{3V} \sum_{i} p_i v_i
$$

$$
P = \frac{1}{3} \int_{0}^{\infty} (p \cdot v) n(p) 4\pi p^2 dp
$$

Blackbody Radiation

$$
n(p) = \frac{1}{h^3} \sum_{j} \frac{g_j}{\exp[(\varepsilon_j + \varepsilon(p) - \mu)/kT] \pm 1}
$$

For photons: $\cdot \quad g = 2$ (2 polarization states)

- $\varepsilon_{\gamma} = 0$ (no excited states)
- $\varepsilon(p) = pc$ (fully relativistic)
- $\mu_{\gamma} = 0$
- bosons, so "-"

$$
n(p) = \frac{1}{h^3} \frac{2}{\exp(pc/kT) - 1}
$$

Blackbody Radiation

Density
\n
$$
n(p) = \frac{1}{h^3} \frac{2}{\exp(pc/kT) - 1}
$$
\n
$$
n_{\gamma} = \int_{0}^{\infty} n(p) 4\pi p^2 dp = \frac{8\pi}{h^3} \int_{0}^{\infty} \frac{p^2 dp}{\exp(pc/kT) - 1}
$$
\n
$$
= \frac{8\pi}{h^3} \left(\frac{kT}{c}\right)^3 \int_{0}^{\infty} \frac{x^2 dx}{e^x - 1} \approx 2.4
$$

$$
= 2.4 \frac{8\pi}{h^3} \left(\frac{kT}{c}\right)^3 = 20.28 T^3 \text{cm}^{-3}
$$

$$
\frac{pc}{kT} = x
$$

$$
dp = \frac{kT}{c}dx
$$

$$
p^2 = \left(\frac{kT}{c}\right)^2 x^2
$$

Blackbody Radiation

Energy

\n
$$
n(p) = \frac{1}{h^3} \frac{2}{\exp(pc/kT) - 1}
$$
\n
$$
E_{\gamma} = \int_{0}^{\infty} (pc) n(p) 4\pi p^2 dp = \frac{8\pi c}{h^3} \int_{0}^{\infty} \frac{p^3 dp}{\exp(pc/kT) - 1}
$$
\n
$$
d p = \frac{kT}{c} dx
$$
\n
$$
= \frac{8\pi c}{h^3} \left(\frac{kT}{c}\right)^4 \int_{0}^{\infty} \frac{x^3 dx}{e^x - 1} = \frac{\pi^4}{15}
$$
\n
$$
= \frac{\pi^4}{15} \frac{8\pi c}{h^3} \left(\frac{kT}{c}\right)^4 = \left(\frac{8\pi^5 k^4}{15h^3 c^3}\right) T^4 \quad \boxed{= aT^4}
$$
\n
$$
a = 7.5 \times 10^{-15} \text{ erg cm}^{-3} \text{K}^{-4}
$$
Blackbody Radiation

$$
n(p) = \frac{1}{h^3} \frac{2}{\exp(pc/kT) - 1}
$$

$$
P_{\gamma} = \frac{1}{3} \int_{0}^{\infty} (pc) n(p) 4\pi p^2 dp = \frac{1}{3} E_{\gamma} \left[= \frac{1}{3} aT^4 \right]
$$

$$
P = (\gamma - 1)E = \frac{1}{3}E \rightarrow \gamma = \frac{4}{3}
$$

Blackbody Radiation

$$
n(p) = \frac{1}{h^3} \frac{2}{\exp(pc/kT) - 1}
$$

The energy density of photons with momentum between *p* and *p+dp* is

$$
\varepsilon(p)n(p)4\pi p^2dp = \frac{8\pi}{h^3} \frac{(pc)p^2dp}{\exp(pc/kT)-1}
$$

The energy density of photons with frequency between *v* and *v+dv* is

$$
\begin{pmatrix} pc = hv \\ dp = (h/c) dv \end{pmatrix} \qquad \frac{8\pi}{h^3} \frac{hv(hv/c)^2 (h/c) dv}{\exp(hv/kT) - 1}
$$

$$
B_v dv = \frac{8\pi h v^3}{c^3} \frac{1}{\exp(hv/kT) - 1} dv
$$

$$
(B_v : \text{erg cm}^{-3} \text{ Hz}^{-1})
$$

Planck function

$$
n(p) = \frac{1}{h^3} \sum_{j} \frac{g_j}{\exp[(\varepsilon_j + \varepsilon(p) - \mu)/kT] \pm 1}
$$

For atoms: $\bullet \ \varepsilon_j = \varepsilon_0$ (single energy state)

•
$$
\varepsilon(p) = \frac{p^2}{2m}
$$
 (non-relativistic)

• $\mu/kT \ll -1$ (+/-1 term can be neglected)

$$
n(p) = \frac{1}{h^3} \frac{g}{\exp[(\varepsilon_0 + p^2/2m - \mu)/kT]}
$$

$$
n(p) = \frac{g}{h^3} e^{\mu/kT} e^{-\varepsilon_0/kT} e^{-p^2/2mkT}
$$

$$
n(p) = \frac{g}{h^3} e^{\mu/kT} e^{-\varepsilon_0/kT} e^{-p^2/2mkT}
$$

Density

=

$$
n = \int_{0}^{\infty} n(p) 4\pi p^{2} dp = \frac{4\pi g}{h^{3}} e^{\mu/kT} e^{-\epsilon_{0}/kT} \int_{0}^{\infty} e^{-p^{2}/2mkT} p^{2} dp
$$

\n
$$
= A \int_{0}^{\infty} e^{-x} (2mkTx) \frac{mkT}{(2mkTx)^{1/2}} dx
$$

\n
$$
= \frac{A}{2} (2mkT)^{3/2} \int_{0}^{\infty} x^{1/2} e^{-x} dx = \frac{\sqrt{\pi}}{2}
$$

\n
$$
p = (2mkT)^{1/2} x^{1/2}
$$

$$
= A \frac{\pi^{1/2}}{4} (2mkT)^{3/2} = \frac{g}{h^3} (2\pi mkT)^{3/2} e^{\mu/kT} e^{-\epsilon_0/kT}
$$

$$
n(p) = \frac{g}{h^3} e^{\mu/kT} e^{-\varepsilon_0/kT} e^{-p^2/2mkT}
$$

$$
P = \frac{1}{3} \int_{0}^{\infty} (p v) n(p) 4 \pi p^{2} dp = \frac{1}{3} \frac{4 \pi g}{h^{3}} e^{\mu/kT} e^{-\epsilon_{0}/kT} \int_{0}^{\infty} \frac{p^{2}}{m} e^{-p^{2}/2mkT} p^{2} dp
$$

$$
=A\frac{1}{3m}\int_{0}^{\infty}e^{-x}\left(2mkTx\right)^{2}\frac{mkT}{\left(2mkTx\right)^{1/2}}dx
$$

 $(2mkT)^{5/2}$ $\int x^{3/2}e^{-x} dx$

∞

0

∫

Pressure

=

1

A

2

3*m*

$$
p^{2}/2mkT = x
$$

$$
dp = \frac{mkT}{p}dx
$$

$$
p = (2mkT)^{1/2}x^{1/2}
$$

$$
= \frac{1}{3m} \frac{A}{2} (2mkT)^{5/2} \left[-x^3 \left(\frac{e^{-x}}{2} \right) \right]_0^{\infty} + \frac{3}{2} \int_0^{\infty} x^{1/2} e^{-x} dx \right]_{v=-e^{-x}}^{u=x^{3/2}} du = (3/2) x^{1/2} dx
$$

$$
= kT \frac{A}{2} (2mkT)^{3/2} \int_{0}^{\infty} x^{1/2} e^{-x} dx \quad \boxed{= nkT}
$$

$$
\text{Energy} \qquad \qquad n(p) = \frac{g}{h^3} e^{\mu/kT} e^{-\epsilon_0/kT} e^{-p^2/2mkT}
$$

$$
E = \int_{0}^{\infty} (p^2/2m)n(p) 4\pi p^2 dp = \frac{3}{2}P \qquad \boxed{\frac{3}{2}nkT}
$$

$$
P = (\gamma - 1)E = \frac{2}{3}E \rightarrow \gamma = \frac{5}{3}
$$

$$
n(p) = \frac{1}{h^3} \sum_{j} \frac{g_j}{\exp[(\varepsilon_j + \varepsilon(p) - \mu)/kT] \pm 1}
$$

For fermions: \bullet $g = 2$ (2 spin states)

•
$$
\varepsilon_0 = mc^2
$$
 (no excited states)

•
$$
\varepsilon(p) = (p^2c^2 + m^2c^4)^{1/2} - mc^2
$$

= $mc^2(\sqrt{1 + (p/mc)^2} - 1)$

• fermions, so "+"

$$
n(p) = \frac{2}{h^3} \frac{1}{\exp\left[\left(\varepsilon(p) - \left(\mu - mc^2\right)\right)/kT\right] + 1}
$$

$$
n(p) = \frac{2}{h^3} \frac{1}{\exp\left[\left(\varepsilon(p) - \left(\mu - mc^2\right)\right)/kT\right] + 1}
$$

A completely degenerate gas behaves as if $T \rightarrow 0$

The probability that an energy state is occupied is

$$
f(\varepsilon) = \frac{1}{\exp\left[\left(\varepsilon(p) - \left(\mu - mc^2\right)\right)/kT\right] + 1} = \begin{cases} 1 & \text{if } \varepsilon < \left(\mu - mc^2\right) \\ 0 & \text{if } \varepsilon > \left(\mu - mc^2\right) \end{cases}
$$

Critical energy: Fermi energy $\varepsilon_F = \mu - mc^2$

Particles cannot have greater energy than this.

Fermi momentum p_F

$$
x = \frac{p}{mc} \rightarrow x_F = \frac{p_F}{mc} \qquad \qquad \varepsilon_F = mc^2 \left(\sqrt{1 + x_F^2} - 1 \right)
$$

Chemical potential is thus $\mu_F = mc^2 + \varepsilon_F$

This is the maximum total energy of particles

$$
n(p) = \frac{2}{h^3} \frac{1}{\exp\left[\left(\varepsilon(p) - \left(\mu - mc^2\right)\right)/kT\right] + 1}
$$

$$
n = \int_{0}^{\infty} n(p) 4\pi p^2 dp = \frac{8\pi}{h^3} \int_{0}^{p_F} p^2 dp = \frac{8\pi}{h^3} \frac{p_F^3}{3}
$$

$$
=\frac{8\pi}{3}\left(\frac{mc}{h}\right)^3 x_F^3
$$

For electrons:
$$
n_e = \frac{8\pi}{3} \left(\frac{m_e c}{h} \right)^3 x_F^3 = 5.9 \times 10^{29} x_F^3
$$
 cm⁻³

 P ressure $n(p)$ = = 8^π $\frac{\partial u}{\partial h^3}\int_0^1 v\cdot p^3 dp$ 0 *pF* ∫ 2 h^3 1 $\exp\left[\left(\varepsilon(p) - \left(\mu - mc^2\right)\right)/kT\right] + 1$ $P =$ 1 $\frac{1}{3}\int_{0}^{1}(p\nu)n(p)4\pi p^{2} dp$ 0 ∞ ∫ $v =$ *d*^ε *dp* $\varepsilon(p) = mc^2 \left(\sqrt{1 + (p/mc)^2} - 1 \right)$ $= mc$ 2 1 2 $1+\left(\frac{p}{p}\right)$ *mc* $\sqrt{}$ ⎝ $\left(\frac{p}{\sqrt{p}}\right)^{n}$ \overline{y} \mathbf{a} $\left(\begin{array}{cc} & \left(& p \end{array}\right)^2$ ⎝ $\overline{}$ \overline{a} \overline{y} and the second $-1/2$ \cdot 2 $\frac{p}{q}$ *mc* $\frac{1}{1}$ *mc* = *p m* $1+\left(\frac{p}{p}\right)$ *mc* $\sqrt{}$ ⎝ $\left(\frac{p}{p}\right)$ ⎠ \mathbf{a} $\left(\begin{array}{cc} p \end{array}\right)^2$ $\overline{\mathsf{L}}$ ⎢ $\overline{}$ $\overline{}$ ⎦ ⎥ $\overline{}$ $-1/2$ = 8^π $3mh³$ *p* 4 *dp* $1 + (p/mc)^2$ 0 *pF* ∫

Pressure

$$
P = \frac{8\pi}{3mh^3} \int_{0}^{p_F} \frac{p^4 dp}{\sqrt{1 + (p/mc)^2}}
$$

$$
\frac{p}{mc} = x
$$

dp = mcdx

$$
p^4 = (mc)^4 x^4
$$

$$
= \frac{8\pi}{3mh^3} \int_0^{x_F} \frac{(mc)^5 x^4 dx}{\sqrt{1+x^2}} = \frac{8\pi m^4 c^5}{3h^3} \int_0^{x_F} \frac{x^4 dx}{\sqrt{1+x^2}}
$$

$$
= \frac{\pi}{3} \left(\frac{mc}{h} \right)^3 mc^2 \left[x_F \left(2x_F^2 - 3 \right) \left(1 + x_F^2 \right)^{1/2} + 3 \sinh^{-1} x_F \right]
$$

$$
\begin{aligned}\n\text{Energy} \quad & \left| n(p) = \frac{2}{h^3} \frac{1}{\exp\left[\left(\varepsilon(p) - \left(\mu - mc^2 \right) \right) / kT \right] + 1} \right| \\
E &= \int_0^\infty \varepsilon(p) n(p) 4\pi p^2 \, dp = \frac{8\pi}{h^3} \int_0^{p_F} mc^2 \left(\sqrt{1 + \left(p/mc \right)^2} - 1 \right) p^2 \, dp \\
&= \frac{8\pi}{h^3} mc^2 \int_0^{x_F} \left(\sqrt{1 + x^2} - 1 \right) (mc)^3 x^2 \, dx \\
&= 8\pi \left(\frac{mc}{h} \right)^3 mc^2 \int_0^{x_F} \left(\sqrt{1 + x^2} - 1 \right) x^2 \, dx \\
&= \frac{p}{m c} = x \\
dp &= mc^2 x^2 \\
\hline\n\end{aligned}
$$

$$
=\frac{\pi}{3}\left(\frac{mc}{h}\right)^3mc^2\left[8x_F^3\left(\sqrt{1+x_F^2}-1\right)-x_F\left(2x_F^2-3\right)\left(1+x_F^2\right)^{1/2}-3\sinh^{-1}x_F\right]
$$

$$
n = \frac{8\pi}{3} \left(\frac{mc}{h}\right)^3 x_F^3 = Ax_F^3
$$

$$
P = \frac{8\pi}{3} \left(\frac{mc}{h}\right)^3 mc^2 \int_0^{x_F} \frac{x^4 dx}{\sqrt{1+x^2}} = Amc^2 \int_0^{x_F} \frac{x^4 dx}{\sqrt{1+x^2}}
$$

$$
E = 8\pi \left(\frac{mc}{h}\right)^3 mc^2 \int_0^{x_F} \left(\sqrt{1 + x^2} - 1\right) x^2 dx = 3Amc^2 \int_0^{x_F} \left(\sqrt{1 + x^2} - 1\right) x^2 dx
$$

$$
n = Ax_F^3
$$

\nLook at limiting
\ncases: NR, UR
\n
$$
P = Amc^2 \int_0^{x_F} \frac{x^4 dx}{\sqrt{1 + x^2}}
$$

\n
$$
E = 3Amc^2 \int_0^{x_F} (\sqrt{1 + x^2} - 1) x^2 dx
$$

$$
\frac{\text{NR:}}{\text{N}} \quad x_F \ll 1 \qquad P \approx A m c^2 \int_0^{x_F} x^4 \, dx = A m c^2 \frac{x_F^5}{5} = A m c^2 \frac{1}{5} \left(\frac{n}{A}\right)^{5/3}
$$
\n
$$
= \frac{m c^2}{5} A^{-2/3} n^{5/3}
$$

$$
E \approx 3Amc^2 \int_0^{x_F} \left(1 + \frac{1}{2}x^2 - 1\right) x^2 dx = \frac{3}{2}Amc^2 \int_0^{x_F} x^4 dx = \frac{3}{2}P
$$

$$
P = (\gamma - 1)E = \frac{2}{3}E \qquad \rightarrow \gamma = \frac{5}{3}
$$

$$
\frac{|n = Ax_F^3|}{\text{Look at limiting}}
$$
\n
$$
P = Amc^2 \int_0^{x_F} \frac{x^4 dx}{\sqrt{1 + x^2}} = \frac{1}{2} \left[E = 3Amc^2 \int_0^{x_F} (\sqrt{1 + x^2} - 1) x^2 dx \right]
$$
\n
$$
Cases: NR, UR
$$

UR:
$$
x_F \gg 1
$$
 $P \approx Amc^2 \int_0^{x_F} x^3 dx = Amc^2 \frac{x_F^4}{4} = Amc^2 \frac{1}{4} \left(\frac{n}{A}\right)^{4/3}$

$$
= \frac{mc^2}{4} A^{-1/3} n^{4/3}
$$

$$
E \approx 3Amc^2 \int_{0}^{x_F} (x-1)x^2 dx = 3Amc^2 \int_{0}^{x_F} x^3 dx = 3P
$$

$$
P = (\gamma - 1)E = \frac{1}{3}E \longrightarrow \gamma = \frac{4}{3}
$$

- gas has same gamma as ideal gas.
- In the relativistic limit, it behaves like radiation.

$$
n = \frac{8\pi}{3} \left(\frac{mc}{h}\right)^3 x_F^3
$$
 Convert to mass density: $n = \frac{N_A \rho}{\mu}$

e.g., electrons:
$$
n_e = \frac{N_A \rho}{\mu_e} = \frac{8\pi}{3} \left(\frac{m_e c}{h}\right)^3 x_F^3
$$

$$
\frac{\rho}{\mu_e} = \frac{8\pi}{3N_A} \left(\frac{m_e c}{h}\right)^3 x_F^3 = 9.7 \times 10^5 x_F^3 \text{ g cm}^{-3}
$$

e.g., neutrons:
$$
n_n = \frac{N_A \rho}{\mu_n} = \frac{8\pi}{3} \left(\frac{m_n c}{h}\right)^3 x_F^3
$$

$$
\frac{\rho}{\mu_n} = \frac{8\pi}{3N_A} \left(\frac{m_n c}{h}\right)^3 x_F^3 = 6.1 \times 10^{15} x_F^3 \text{ g cm}^{-3}
$$

Useful approximations

• Partly degenerate gas

$$
P_e \approx \left(P_{e,nd}^2 + P_{e,d}^2\right)^{1/2}
$$

• Partly relativistic gas

$$
P_{e,d} \approx \left(P_{e,d,NR}^{-2} + P_{e,d,UR}^{-2}\right)^{-1/2}
$$

This formula picks out the smallest pressure

These interpolation formulae are good to ~2%

Equation of State Summary

- Start with ρ , T , X_i at some point in star
- Assume Local Thermodynamic Equilibrium
- Total gas pressure is the sum of components

$$
P = P_{\text{rad}} + P_{\text{ion}} + P_e
$$

• The radiation pressure is

$$
P_{\text{rad}} = \frac{1}{3} aT^4 \left(a = 7.5 \times 10^{-15} \text{ erg cm}^{-3} \text{K}^{-4} \right)
$$

• The ion ideal gas pressure is

$$
P_{\text{ion}} = \frac{N_A k}{\mu_I} \rho T \qquad (N_A = 6.022 \times 10^{23} \text{ mole}^{-1})
$$

Equation of State Summary

• The electron pressure can be a mixture of non-degenerate and degenerate pressure

$$
P_e \approx \left(P_{e,nd}^2 + P_{e,d}^2\right)^{1/2}
$$

• Where the electron non-degenerate pressure is

$$
P_{e,nd} = \frac{N_A k}{\mu_e} \rho T
$$

• And the electron degenerate pressure can be non-relativistic or relativistic

$$
\boxed{P_{e,d} \approx \left(P_{e,d,NR}^{-2} + P_{e,d,UR}^{-2}\right)^{-1/2}}
$$

Equation of State Summary

• Where the non-relativistic, degenerate pressure is

$$
P_{e,d,NR} = \left(\frac{3}{8\pi}\right)^{2/3} \frac{h^2}{5m_e} \left(\frac{N_A}{\mu_e}\right)^{5/3} \rho^{5/3}
$$

• And the ultra-relativistic, degenerate pressure is

$$
P_{e,d,UR} = \left(\frac{3}{8\pi}\right)^{1/3} \frac{hc}{4} \left(\frac{N_A}{\mu_e}\right)^{4/3} \rho^{4/3}
$$

• Compute the mean molecular weights from *Xi*

$$
\mu_i^{-1} = \sum_i \frac{X_i}{A_i} \qquad \mu_e^{-1} = \sum_i \frac{X_i Z_i y_i}{A_i}
$$

• Get the ionization fractions from the Saha equation.

The Boltzmann Equation

Ideal gas :
$$
n = g \left(\frac{2\pi mkT}{h^2} \right)^{3/2} e^{\mu/kT} e^{-\epsilon_0/kT}
$$

Consider two species of particles: atoms in different energy levels that can be excited or de-excited via photons.

e.g.,
$$
\frac{H_1 + \gamma \rightarrow H_2}{n_1} = \frac{g_1 (2\pi m k \chi/h^2)^{3/2} e^{\chi/kT} e^{-\epsilon_1/kT}}{g_2 (2\pi m k T/h^2)^{3/2} e^{\mu_2/kT} e^{-\epsilon_2/kT}} = \frac{g_1 e^{-\epsilon_1/kT}}{g_2 e^{-\epsilon_2/kT}}
$$

 $g₂$

 $n₂$

The Boltzmann Equation

Total number of atoms:
$$
n_{\text{tot}} = \sum_{i} n_i
$$

 n_{tot} n_i *n_i* n_i *g_i* = $n₁$ + $n₂$ $+ \cdots$ = *g*1 $e^{-\varepsilon_1/kT}e^{\varepsilon_i/kT} + \frac{g_2}{2}e^{-\varepsilon_2/kT}e^{\varepsilon_i/kT} + \cdots$ *gi*

$$
= \frac{1}{g_i} e^{\varepsilon_i/kT} \left(g_1 e^{-\varepsilon_1/kT} + g_2 e^{-\varepsilon_2/kT} + \cdots \right) = \frac{1}{g_i e^{-\varepsilon_i/kT}} \sum_j g_j e^{-\varepsilon_j/kT}
$$

$$
\frac{n_i}{n_{\text{tot}}} = \frac{g_i}{G} e^{-\varepsilon_i / kT}
$$

Partition function:
$$
G = \sum_{j} g_j e^{-\varepsilon_j / kT}
$$

Like an effective statistical weight for all states

Consider an atomic gas that is partly ionized

e.g., $H^+ + e^- \leftrightarrow H^0 + \gamma \Big| 1\mu^+ + 1\mu^- = 1\mu^0$

$$
\varepsilon_{+} = \chi
$$

$$
\varepsilon_{0} = 0
$$

• charged ions in ground state $n^+ = g$

$$
g^{+}\left(\frac{2\pi m_{p}kT}{h^{2}}\right)^{3/2}e^{\mu^{+}/kT}e^{-\chi/kT}
$$

• free electrons

$$
n^{-} = g^{-}\left(\frac{2\pi m_e kT}{h^2}\right)^{3/2} e^{\mu^{-}/kT}
$$

• neutral ions in ground state

$$
n^{0}=g^{0}\left(\frac{2\pi m_{p}kT}{h^{2}}\right)^{3/2}e^{\mu^{0}/kT}
$$

Combine to eliminate chemical potentials

$$
\frac{n^+n^-}{n^0} = \frac{g^+g^-}{g^0} \left(\frac{2\pi m_e kT}{h^2}\right)^{3/2} e^{-\chi/kT}
$$

$$
\frac{n^+n^-}{n^0} = \frac{g^+g^-}{g^0} \left(\frac{2\pi m_e kT}{h^2}\right)^{3/2} e^{-\chi/kT}
$$

This is only for atoms in the ground state.

In general, we have neutral and ionized atoms in multiple states Use Boltzmann to generalize to all states of a given atom

$$
\frac{n^{\dagger}n_{e}}{n^{0}} = \frac{2G^{+}}{G^{0}} \left(\frac{2\pi m_{e}kT}{h^{2}}\right)^{3/2} e^{-\chi/kT}
$$

Example: pure H

$$
n^{+} = yn_{I}
$$

\n
$$
n^{0} = (1 - y)n_{I}
$$

\n
$$
n_{e} = yn_{I}
$$

\n
$$
\frac{p^{+}n_{e}}{n^{0}} = \frac{yn_{I} \cdot yn_{I}}{(1 - y)n_{I}} = \frac{y^{2}}{1 - y}n_{I}
$$

\n
$$
\frac{y^{2}}{1 - y} = \frac{1}{n_{I}} \left(\frac{2\pi m_{e}kT}{h^{2}}\right)^{3/2} e^{-\chi/kT} n_{I} = N_{A}\rho
$$

y 2 1− *y* = 4×10^{-9} g cm⁻³ ρ $T^{3/2}e^{-1.578\times10^5/T}$ 50% ionized at T=10⁴K for low densities

Example: Balmer line strength Strength of Balmer lines depend on fraction of H gas that is in the n=2 excited state.

 Ω

 Ω

 Ω

13.6eV
\n
$$
n = \infty
$$
\n10.2eV
\n
\n
$$
n = 3
$$
\n
$$
n = 2
$$

 n_2^0 *nI* $n=1$ 0 We want n_2^0 $\frac{n_2}{n^0}$ = g_2^0 Boltzmann gives us $\frac{n_2}{n^0} = \frac{82}{G^0} e^{-\epsilon_2/kT}$

$$
\frac{n_2^0}{n_1} = \frac{n_2^0}{n_1^0} \frac{n_1^0}{n_1}
$$
\n
$$
\frac{n_2^0}{n_1} = (1 - y) \frac{g_2^0}{G^0} e^{-\epsilon_2/kT}
$$
\n
$$
\frac{g_2^0 = 8}{G^0} = \sum_i g_i e^{-\epsilon_i/kT} \approx 2
$$
\n
$$
\epsilon_2 = 10.2 \text{ eV}
$$

$$
\frac{n_2^0}{n^0} = (1-y)\frac{g_2^0}{G^0}e^{-\epsilon_2/kT}
$$

$$
\frac{n_2^0}{n^0} = (1-y)\frac{g_2^0}{G^0}e^{-\epsilon_2/kT}
$$
\n
$$
\frac{y^2}{1-y} = \frac{1}{n}\left(\frac{2\pi m_e kT}{h^2}\right)^{3/2}e^{-\chi/kT}
$$

A different way to write the Saha equation

$$
\frac{n^{\dagger}n_e}{n^0} = \frac{2G^{\dagger}}{G^0} \left(\frac{2\pi m_e kT}{h^2}\right)^{3/2} e^{-\chi/kT}
$$

$$
\log\left(\frac{n^+n_e}{n^0}\right) = \log\left(\frac{2G^+}{G^0}\right) + \frac{3}{2}\log\left(\frac{2\pi m_e kT}{h^2}\right) + \log\left(e^{-\chi/kT}\right)
$$

$$
\log\left(\frac{n^+}{n^0}\right) + \log n_e = \log\left(\frac{2G^+}{G^0}\right) + \frac{3}{2}\log\left(\frac{2\pi m_e k}{h^2}\right) + \frac{3}{2}\log T - \ln 10\frac{\chi}{kT}
$$

$$
P_e = n_e kT \rightarrow \log n_e = \log P_e - \log k - \log T
$$

$$
\log\left(\frac{n^+}{n^0}\right) = \log\left(\frac{2G^+}{G^0}\right) + \frac{5}{2}\log T - \log P_e - \frac{5040\chi}{T} - 0.48
$$

The Saha equation works best at moderate densities

- At very low densities (e.g., corona), we must worry about whether LTE applies.
- At very high densities, atoms are not isolated.

Overlap of potentials of neighboring atoms lowers the effective ionization energy, which leads to greater ionization than Saha predicts.

For density *n* and separation $a : n =$ 4 3 $\int \frac{4}{\pi} \pi a^3$ ⎝ $\left(\frac{4}{3}\pi a^3\right)^3$ ⎠ ⎟ −1

set *a* = radius of first Bohr orbit of $H = 0.5 \times 10^{-8}$ cm $\rightarrow \rho \sim 3$ g cm⁻³

Above this density, even the ground state is affected and all H is ionized.

• For ions of charge Z, the ratio of electrostatic energy to thermal energy is a measure of whether Coulomb effects affect the ideal-ness of an ideal gas.

$$
\Gamma_C = \frac{Z^2 e^2 / a}{kT} \qquad \qquad \Gamma_C = 1 \to \rho = 85 \text{ g cm}^{-3} \left(\frac{T}{10^6 K}\right)^3
$$

Important in very low mass stars

Equation of State

For pure H

- ideal gas vs. radiation
-
- $ρ = 1.5 × 10⁻²³T³$

 degenerate vs. not
 $ρ = 6 × 10⁻⁹T^{3/2}$

 neutral vs. ionized
 $ρ = 8 × 10⁻⁹T^{3/2}e^{-1.58}$

 pressure ionized
 $ρ ≈ 1$

 Coulomb effects
 $ρ = 8.5 × 10⁻¹⁷T³$

 $\times 10^5\big/ T$
- $\rho \approx 1$
- $\rho = 8.5 \times 10^{-17} T^3$
- $\rho = 4.2 \times 10^{-10} T^3$

Adiabatic Exponents

- *Q* : specific heat $(\text{erg } g^{-1})$
- V_ρ : specific volume $\left(\text{cm}^3 \text{g}^{-1}\right)$ = 1 ρ
- *E* : specific internal energy $(\text{erg } g^{-1})$

 $dQ = dE + PdV_{\rho}$ = $dE + Pd\left(\frac{1}{2}\right)$ ρ \int ⎝ $\overline{}$ \overline{a} ⎠ $\int dE - \frac{P}{\rho^2} d\rho$ First law of thermodynamics $\Gamma_1 =$ ∂ln*P* $\partial \ln \rho$ \int ⎝ $\overline{}$ \overline{a} ⎠ ⎟ ad $=-\frac{\partial \ln P}{\partial \ln P}$ ∂ln*V*^ρ \int ⎝ $\mathsf L$ \overline{a} ⎠ and the same of ad Γ_{2} Γ_2 – 1 = ∂ln*P* ∂ln*T* $\big($ ⎝ $\left(\frac{\partial \ln P}{\partial \ln T}\right)^n$ ⎠ ⎟ ad = 1 ∇_{ad} $\Gamma_3 - 1 =$ ∂ln*T* $\partial \ln \rho$ $\big($ ⎝ $\overline{}$ $\overline{}$ ⎠ ⎟ ad $=-\frac{\partial \ln T}{\partial \ln T}$ ∂ln*V*^ρ \int ⎝ $\mathsf L$ \overline{a} ⎠ and the same of ad $\nabla_{_{\mathrm{ad}}}=$ ∂ln*T* $\partial \ln \rho$ $\big($ ⎝ $\overline{}$ \overline{a} \overline{y} \mathbf{a} ad $\partial \ln \rho$ ∂ln*P* $\big($ ⎝ $\left(\frac{\partial \ln \rho}{\partial \ln \rho}\right)^n$ \overline{y} \mathbf{a} ad = Γ_3 – 1 Γ_{1} = Γ_2 – 1 Γ_{2}

Adiabatic Exponents

- \cdot Γ ₁ describes how pressure responds to compression (relevant for dynamical processes like pulsations)
- Γ ₂ describes how temperature responds to changes in pressure (relevant for determining whether convection takes place)
- \cdot Γ ₃ describes how temperature responds to compression

Ideal Gas:	\n $\Gamma_1 = \Gamma_2 = \Gamma_3 = \frac{5}{3}$ \n	\n Γ_3 is equivalent to the γ in the γ -law equation of state
Radiation:	\n $\Gamma_1 = \Gamma_2 = \Gamma_3 = \frac{4}{3}$ \n	\n $P = (\gamma - 1)\rho E$ \n

For mixtures of gas and radiation, as well as for cases where chemical reaction happen, the adiabatic exponents can differ from each other.
Adiabatic Exponents: Methodology

$$
P = f(T, V_{\rho})
$$

$$
E
$$

$$
E = g(T, V_{\rho})
$$

Adiabatic change $dQ = 0$

$$
dQ = dE + PdV_{\rho} = 0
$$

= $\left(\frac{\partial E}{\partial T}\right)_{V_{\rho}} dT + \left(\frac{\partial E}{\partial V_{\rho}}\right)_{T} dV_{\rho} + PdV_{\rho} = 0$

$$
dP = \left(\frac{\partial P}{\partial T}\right)_{V_{\rho}} dT + \left(\frac{\partial P}{\partial V_{\rho}}\right)_{T} dV_{\rho} \qquad = \left(\frac{\partial P}{\partial T}\right)_{ad} dT = \left(\frac{\partial P}{\partial V_{\rho}}\right)_{ad} dV_{\rho}
$$

$$
= \left(\frac{\Gamma_2}{\Gamma_2 - 1}\right) dT = (-\Gamma_1) dV_\rho
$$

Pressure

$$
P = P_{\rm gas} + P_{\rm rad} = \frac{N_A k}{\mu} \rho T + \frac{1}{3} a T^4 \qquad \qquad = \frac{N_A k}{\mu} \frac{T}{V_{\rho}} + \frac{1}{3} a T^4
$$

Specific Internal Energy

$$
E = E_{\rm gas} + E_{\rm rad} = \frac{3}{2} \frac{N_A k}{\mu} T + a \frac{T^4}{\rho} \qquad \left| = \frac{3}{2} \frac{N_A k}{\mu} T + a T^4 V_{\rho} \right|
$$

$$
P = \frac{N_A k}{\mu} \frac{T}{V_\rho} + \frac{1}{3} aT^4
$$

$$
aT^4 \qquad E = \frac{3}{2} \frac{N_A k}{\mu} T + aT^4 V_\rho
$$

$$
\begin{array}{c}\n \text{Adiabatic change} \\
 dQ = 0\n \end{array}
$$

$$
dQ = \left(\frac{\partial E}{\partial T}\right)_V dT + \left(\frac{\partial E}{\partial V_\rho}\right)_T dV_\rho + P dV_\rho = 0
$$

$$
= \left(\frac{3N_A k}{2\mu} + 4aT^3 V_\rho\right) dT + \left(aT^4\right) dV_\rho + \left(\frac{N_A k}{\mu} \frac{T}{V_\rho} + \frac{1}{3} aT^4\right) dV_\rho = 0
$$

$$
= \left(\frac{3N_A k}{2\mu} + 4aT^3 V_\rho\right) dT + \left(\frac{N_A k}{\mu} \frac{T}{V_\rho} + \frac{4}{3} aT^4\right) dV_\rho = 0
$$

$$
= \left(\frac{3N_A k}{2\mu} \frac{T}{V_\rho} + 4aT^4\right) \frac{V_\rho}{T} dT + \left(\frac{N_A k}{\mu} \frac{T}{V_\rho} + \frac{4}{3} aT^4\right) dV_\rho = 0
$$

$$
=\left(\frac{3}{2}P_{\text{gas}} + 12P_{\text{rad}}\right)\frac{dT}{T} + \left(P_{\text{gas}} + 4P_{\text{rad}}\right)\frac{dV_{\rho}}{V_{\rho}} = 0
$$

$$
P = \frac{N_A k}{\mu} \frac{T}{V_\rho} + \frac{1}{3} aT^4
$$

$$
aT^4 \qquad \qquad E = \frac{3}{2} \frac{N_A k}{\mu} T + aT^4 V_\rho
$$

Equation of state

$$
dP = \left(\frac{\partial P}{\partial T}\right)_{V_{\rho}} dT + \left(\frac{\partial P}{\partial V_{\rho}}\right)_{T} dV_{\rho} = \left(\frac{N_{A}k}{\mu} \frac{1}{V_{\rho}} + \frac{4}{3} aT^{3}\right) dT + \left(-\frac{N_{A}k}{\mu} \frac{T}{V_{\rho}^{2}}\right) dV_{\rho}
$$

$$
= \left(\frac{N_{A}k}{\mu} \frac{T}{V_{\rho}} + \frac{4}{3} aT^{4}\right) \frac{1}{T} dT + \left(-\frac{N_{A}k}{\mu} \frac{T}{V_{\rho}}\right) \frac{1}{V_{\rho}} dV_{\rho}
$$

$$
= \left(P_{\text{gas}} + 4P_{\text{rad}}\right) \frac{dT}{T} - P_{\text{gas}} \frac{dV_{\rho}}{V_{\rho}} = dP = \left(\frac{\partial P}{\partial T}\right)_{\text{ad}} dT = \frac{P}{T} \left(\frac{\partial \ln P}{\partial \ln T}\right)_{\text{ad}} dT
$$

$$
\rightarrow \left(P_{\text{gas}} + 4P_{\text{rad}}\right)\frac{dT}{T} - P_{\text{gas}}\frac{dV_{\rho}}{V_{\rho}} - \frac{\Gamma_2}{\Gamma_2 - 1}\left(P_{\text{gas}} + P_{\text{rad}}\right)\frac{dT}{T} = 0
$$

$$
= \left(P_{\text{gas}} + 4P_{\text{rad}} - \frac{\Gamma_2}{\Gamma_2 - 1} \left(P_{\text{gas}} + P_{\text{rad}}\right)\right) \frac{dT}{T} - P_{\text{gas}} \frac{dV_{\rho}}{V_{\rho}} = 0
$$

1st Law of Thermodynamics	$\left(\frac{3}{2}P_{\text{gas}} + 12P_{\text{rad}}\right)\frac{dT}{T} + \left(P_{\text{gas}} + 4P_{\text{rad}}\right)\frac{dV_{\rho}}{V_{\rho}} = 0$		
Equation of State	$\left(P_{\text{gas}} + 4P_{\text{rad}} - \frac{\Gamma_{2}}{\Gamma_{2} - 1}\left(P_{\text{gas}} + P_{\text{rad}}\right)\right)\frac{dT}{T} - P_{\text{gas}}\frac{dV_{\rho}}{V_{\rho}} = 0$		
$Ax + By = 0$	$\rightarrow \frac{A}{C} = \frac{B}{D}$	$\frac{P_{\text{gas}} + 4P_{\text{rad}} - \frac{\Gamma_{2}}{\Gamma_{2} - 1}\left(P_{\text{gas}} + P_{\text{rad}}\right)}{\frac{3}{2}P_{\text{gas}} + 12P_{\text{rad}}}$	$\frac{P_{\text{gas}} + 4P_{\text{rad}}}{P_{\text{gas}} + 4P_{\text{rad}}}$
$P_{\text{gas}} = \beta P$	$\beta R + 4\left(1 - \beta\right)R - \frac{\Gamma_{2}}{\Gamma_{2} - 1}R$	βR	
$P_{\text{gas}} = (1 - B)P$	βR		

3

2

 $\frac{1}{2} - 1$

 $\beta P + 4(1-\beta)P$

 βR + 12(1 – β)*R*

$$
=(1-\beta)P
$$

 $P_{\rm rad}$ ^{$\,$}

$$
\frac{\beta + 4(1-\beta) - \frac{\Gamma_2}{\Gamma_2 - 1}}{2\beta + 12(1-\beta)} = -\frac{\beta}{\beta + 4(1-\beta)} \quad \to \quad \frac{4 - 3\beta - \frac{\Gamma_2}{\Gamma_2 - 1}}{12 - \frac{21}{2}\beta} = -\frac{\beta}{4 - 3\beta}
$$

$$
\rightarrow \left(4 - 3\beta - \frac{\Gamma_2}{\Gamma_2 - 1}\right) \left(4 - 3\beta\right) = \left(-\beta\right) \left(12 - \frac{21}{2}\beta\right)
$$

$$
\rightarrow 16 - 12\beta - 12\beta + 9\beta^2 - \frac{\Gamma_2}{\Gamma_2 - 1}(4 - 3\beta) = -12\beta + \frac{21}{2}\beta^2
$$

$$
\rightarrow \frac{\Gamma_2}{\Gamma_2 - 1} (4 - 3\beta) = 16 - 12\beta - \frac{3}{2} \beta^2 \rightarrow \frac{\Gamma_2 - 1}{\Gamma_2} = \frac{4 - 3\beta}{16 - 12\beta - \frac{3}{2} \beta^2}
$$

$$
\rightarrow \frac{1}{\Gamma_2} = 1 - \frac{4 - 3\beta}{16 - 12\beta - \frac{3}{2}\beta^2} = \frac{16 - 12\beta - \frac{3}{2}\beta^2 - 4 + 3\beta}{16 - 12\beta - \frac{3}{2}\beta^2} = \frac{12 - 9\beta - \frac{3}{2}\beta^2}{16 - 12\beta - \frac{3}{2}\beta^2}
$$

$$
\frac{1}{\Gamma_2} = \frac{12 - 9\beta - \frac{3}{2}\beta^2}{16 - 12\beta - \frac{3}{2}\beta^2} \quad \to \Gamma_2 = \frac{16 - 12\beta - \frac{3}{2}\beta^2}{12 - 9\beta - \frac{3}{2}\beta^2} \quad \to \Gamma_2 = \frac{32 - 24\beta - 3\beta^2}{24 - 18\beta - 3\beta^2}
$$

Similarly, we get:

$$
\Gamma_1 = \frac{32 - 24\beta - 3\beta^2}{24 - 21\beta}
$$

$$
\Gamma_2 = \frac{32 - 24\beta - 3\beta^2}{24 - 18\beta - 3\beta^2}
$$

$$
\Gamma_3 = \frac{32 - 27\beta}{24 - 21\beta}
$$

Thermodynamics of partially ionized gas is different because

- The number of free particles is not constant
- Ionization energy is required to increase n

Assuming a pure H ideal gas

$$
n^{+} = yn_{I}
$$

\n
$$
n^{0} = (1 - y)n_{I}
$$

\n
$$
n_{e} = yn_{I}
$$

\n
$$
n_{I} = N_{A}\rho
$$

\n
$$
n_{I} = \frac{N_{A}\rho}{\sqrt{N_{A}\rho}} = \frac{1}{N_{A}\rho} \left(\frac{2\pi m_{e}kT}{h^{2}}\right)^{3/2} e^{-\chi/kT}
$$

$$
\frac{y^2}{1-y} = \frac{A}{N_A} V_{\rho} T^{3/2} e^{-\chi/kT}
$$

Pressure

$$
P = (n0 + n+ + ne)kT = [(1 - y)nI + ynI + ynI]kT
$$

$$
= (1+y)n_{I}kT = (1+y)N_{A}\rho kT \rightarrow \left| P = (1+y)N_{A}k \frac{T}{V_{\rho}}
$$

Specific Internal Energy

$$
E = \frac{3}{2} (n^{0} + n^{+} + n_{e}) \frac{kT}{\rho} + \frac{n^{+} \chi}{\rho} = (1 + y) n_{I} \frac{3}{2} \frac{kT}{\rho} + \frac{yn_{I}}{\rho} \chi
$$

$$
E = (1 + y)\frac{3}{2}N_A kT + yN_A \chi
$$

$$
P = (1+y)N_A k \frac{T}{V_{\rho}} \left[E = (1+y)\frac{3}{2}N_A kT + yN_A \chi \right]
$$
 Adiabatic chi

$$
\begin{array}{|l|}\n\hline\n\text{Adiabatic change} \\
\hline\n\text{d}Q = 0\n\end{array}
$$

$$
dQ = \left(\frac{\partial E}{\partial T}\right)_{V_{\rho, y}} dT + \left(\frac{\partial E}{\partial V_{\rho}}\right)_{T, y} dV_{\rho} + \left(\frac{\partial E}{\partial y}\right)_{T, V_{\rho}} dy + PdV_{\rho} = 0
$$

\n
$$
= \left[(1+y)\frac{3}{2}N_{A}k \right] dT + \left[\frac{3}{2}N_{A}kT + N_{A}\chi \right] dy + \left[(1+y)N_{A}k\frac{T}{V_{\rho}} \right] dV_{\rho} = 0
$$

\n
$$
= \left[(1+y)\frac{3}{2}N_{A}kT \right] \frac{dT}{T} + \left[(1+y)\frac{3}{2}N_{A}kT + (1+y)N_{A}\chi \right] \frac{dy}{1+y}
$$

\n
$$
+ \left[(1+y)N_{A}kT \right] \frac{dV_{\rho}}{V_{\rho}} = 0
$$

\n
$$
= E_{\text{ideal}} \left[\frac{dT}{T} + \frac{2}{3} \frac{dV_{\rho}}{V_{\rho}} + \left(1 + \frac{2}{3} \frac{\chi}{kT} \right) \frac{dy}{1+y} \right] = 0
$$

$$
P = (1+y)N_A k \frac{T}{V_{\rho}} \left[E = (1+y)\frac{3}{2}N_A kT + yN_A \chi \right]
$$
 Equation

Equation
of State

$$
dP = \left(\frac{\partial P}{\partial T}\right)_{V_{\rho, y}} dT + \left(\frac{\partial P}{\partial V_{\rho}}\right)_{T, y} dV_{\rho} + \left(\frac{\partial P}{\partial y}\right)_{T, V_{\rho}} dy
$$

$$
= \left[(1+y)N_A k \frac{1}{V_{\rho}} \right] dT - \left[(1+y)N_A k \frac{T}{V_{\rho}^2} \right] dV_{\rho} + \left[N_A k \frac{T}{V_{\rho}} \right] dy
$$

$$
= \left[(1+y)N_A k \frac{T}{V_{\rho}} \right] \frac{dT}{T} - \left[(1+y)N_A k \frac{T}{V_{\rho}} \right] \frac{dV_{\rho}}{V_{\rho}} + \left[(1+y)N_A k \frac{T}{V_{\rho}} \right] \frac{dy}{1+y}
$$

$$
dP = (1+y)N_A k \frac{T}{V_{\rho}} \left[\frac{dT}{T} - \frac{dV_{\rho}}{V_{\rho}} + \frac{dy}{1+y} \right] \rightarrow \quad dP = P \left[\frac{dT}{T} - \frac{dV_{\rho}}{V_{\rho}} + \frac{dy}{1+y} \right]
$$

$$
f(y) = \frac{y^2}{1-y} = \frac{A}{N_A} V_{\rho} T^{3/2} e^{-\chi/kT}
$$

Saha Equation

$$
dy = \frac{dy}{df} df = \frac{dy}{df} \left[\left(\frac{\partial f}{\partial T} \right)_{V_{\rho}} dT + \left(\frac{\partial f}{\partial V_{\rho}} \right)_{T} dV_{\rho} \right]
$$

$$
= \frac{(1-y)^2}{2y(1-y)+y^2} \left[\frac{A}{N_A} V_\rho \left(\frac{3}{2} T^{1/2} e^{-\chi/kT} + T^{3/2} e^{-\chi/kT} \frac{\chi}{kT^2} \right) dT + \frac{A}{N_A} T^{3/2} e^{-\chi/kT} dV_\rho \right]
$$

$$
= \frac{(1-y)^2}{2y(1-y)+y^2} \frac{A}{N_A} V_\rho T^{3/2} e^{-\chi/kT} \left[\left(\frac{3}{2} \frac{1}{T} + \frac{\chi}{kT^2} \right) dT + \frac{dV_\rho}{V_\rho} \right]
$$

$$
= \frac{(1-y)^2}{2y(1-y) + y^2} \frac{y^2}{1-y} \left[\left(\frac{3}{2} + \frac{\chi}{kT} \right) \frac{dT}{T} + \frac{dV_{\rho}}{V_{\rho}} \right]
$$

 $\overline{}$

 $\frac{1}{2}$ ⎥

 $\overline{\mathsf{I}}$

$$
dy = \frac{(1-y)^2}{2y(1-y)+y^2} \frac{y^2}{1-y} \left[\left(\frac{3}{2} + \frac{\chi}{kT} \right) \frac{dT}{T} + \frac{dV_{\rho}}{V_{\rho}} \right]
$$

$$
dy = \frac{(1-y)y}{2(1-y)+y} \left[\left(\frac{3}{2} + \frac{\chi}{kT} \right) \frac{dT}{T} + \frac{dV_{\rho}}{V_{\rho}} \right]
$$

$$
dy = \frac{(1-y)y}{2-y} \left[\left(\frac{3}{2} + \frac{\chi}{kT} \right) \frac{dT}{T} + \frac{dV_{\rho}}{V_{\rho}} \right]
$$

$$
\frac{dy}{1+y} = \frac{(1-y)y}{(1+y)(2-y)} \left[\left(\frac{3}{2} + \frac{\chi}{kT} \right) \frac{dT}{T} + \frac{dV_{\rho}}{V_{\rho}} \right]
$$

1st Law of **Thermodynamics**

$$
E_{\text{ideal}} \left[\frac{dT}{T} + \frac{2}{3} \frac{dV_{\rho}}{V_{\rho}} + \left(1 + \frac{2}{3} \frac{\chi}{kT} \right) \frac{dy}{1+y} \right] = 0
$$

Equation of State

$$
dP = P\left[\frac{dT}{T} - \frac{dV_{\rho}}{V_{\rho}} + \frac{dy}{1+y}\right]
$$

Saha

$$
\frac{dy}{1+y} = D(y) \left[\left(\frac{3}{2} + \frac{\chi}{kT} \right) \frac{dT}{T} + \frac{dV_{\rho}}{V_{\rho}} \right]
$$

1st Law of **Thermodynamics**

$$
E_{\text{ideal}} \left[\frac{dT}{T} + \frac{2}{3} \frac{dV_{\rho}}{V_{\rho}} + \left(1 + \frac{2 \chi}{3 kT} \right) \frac{dy}{1+y} \right] = 0
$$

$$
E_{\text{ideal}} \left[\frac{dT}{T} + \frac{2}{3} \frac{dV_{\rho}}{V_{\rho}} + \left(1 + \frac{2 \chi}{3 kT} \right) D(y) \left[\left(\frac{3}{2} + \frac{\chi}{kT} \right) \frac{dT}{T} + \frac{dV_{\rho}}{V_{\rho}} \right] \right] = 0
$$

$$
E_{\text{ideal}} \left[\left[1 + \left(1 + \frac{2 \chi}{3 kT} \right) D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right) \right] \frac{dT}{T} + \left[\frac{2}{3} + \left(1 + \frac{2 \chi}{3 kT} \right) D(y) \right] \frac{dV_{\rho}}{V_{\rho}} \right] = 0
$$

$$
E_{\text{ideal}} \left[\left[1 + D(y) \frac{2}{3} \left(\frac{3}{2} + \frac{\chi}{kT} \right)^2 \right] \frac{dT}{T} + \frac{2}{3} \left[1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right) \right] \frac{dV_{\rho}}{V_{\rho}} \right] = 0
$$

Equation of State

$$
dP = P \left[\frac{dT}{T} - \frac{dV_{\rho}}{V_{\rho}} + \frac{dy}{1+y} \right]
$$

$$
dP = P \left[\frac{dT}{T} - \frac{dV_{\rho}}{V_{\rho}} + D(y) \left[\left(\frac{3}{2} + \frac{\chi}{kT} \right) \frac{dT}{T} + \frac{dV_{\rho}}{V_{\rho}} \right] \right]
$$

$$
dP = P \left[\left[1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right) \right] \frac{dT}{T} + \left[D(y) - 1 \right] \frac{dV_{\rho}}{V_{\rho}} \right]
$$

$$
dP = \left(\frac{\partial P}{\partial T}\right)_{\text{ad}} dT = \frac{P}{T} \left(\frac{\partial \ln P}{\partial \ln T}\right)_{\text{ad}} dT = \frac{P}{T} \frac{\Gamma_2}{\Gamma_2 - 1} dT
$$

 Γ_{2} Γ_2 – 1 *dT T* $=\left(1+ D(y) \right) \frac{3}{2}$ 2 $+\frac{\chi}{4}$ *kT* $\sqrt{}$ ⎝ $\left(\frac{3}{2} + \frac{\chi}{kT}\right)$ ⎠ $1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right)$ \lfloor $\left|1+D(y)\left(\frac{3}{2}+\frac{\chi}{kT}\right)\right|$ $\overline{}$ \vert *dT T* $+\lfloor D(y)-1\rfloor$ *dV*^ρ *V*ρ

$$
\left[1 - \frac{\Gamma_2}{\Gamma_2 - 1} + D(y)\left(\frac{3}{2} + \frac{\chi}{kT}\right)\right] \frac{dT}{T} + \left[D(y) - 1\right] \frac{dV_\rho}{V_\rho} = 0
$$

$$
\frac{\mathbf{1^{st} Law}}{\mathbf{1^{st} Law}} \left[1 + D(y) \frac{2}{3} \left(\frac{3}{2} + \frac{\chi}{kT} \right)^2 \right] \frac{dT}{T} + \frac{2}{3} \left[1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right) \right] \frac{dV_{\rho}}{V_{\rho}} = 0
$$
\n
$$
\frac{\text{Equation of State}}{\text{of State}} \left[1 - \frac{\Gamma_2}{\Gamma_2 - 1} + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right) \right] \frac{dT}{T} + \left[D(y) - 1 \right] \frac{dV_{\rho}}{V_{\rho}} = 0
$$

$$
\frac{1 - \frac{\Gamma_2}{\Gamma_2 - 1} + D(y) \left(\frac{3}{2} + \frac{\chi}{kT}\right)}{1 + D(y) \frac{2}{3} \left(\frac{3}{2} + \frac{\chi}{kT}\right)^2} = \frac{D(y) - 1}{\frac{2}{3} \left[1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT}\right)\right]}
$$

$$
\frac{\Gamma_2}{\Gamma_2 - 1} = 1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right) - \frac{\left(D(y) - 1 \right) \left[1 + D(y) \frac{2}{3} \left(\frac{3}{2} + \frac{\chi}{kT} \right)^2 \right]}{\frac{2}{3} \left[1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right) \right]}
$$

$$
\frac{\Gamma_2}{\Gamma_2 - 1} = 1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right) - \frac{\left(D(y) - 1 \right) \left[1 + D(y) \frac{2}{3} \left(\frac{3}{2} + \frac{\chi}{kT} \right)^2 \right]}{\frac{2}{3} \left[1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right) \right]}
$$

…algebra…

$$
\frac{\Gamma_2}{\Gamma_2 - 1} = \frac{\frac{5}{2} + D(y) \left[\frac{15}{4} + 5 \frac{\chi}{kT} + \left(\frac{\chi}{kT} \right)^2 \right]}{1 + D(y) \left(\frac{3}{2} + \frac{\chi}{kT} \right)}
$$

$$
D(y) = \frac{y(1 - y)}{(1 + y)(2 - y)}
$$

Similarly:

$$
\Gamma_3 - 1 = \frac{2 + 2D(y) \left(\frac{3}{2} + \frac{\chi}{kT}\right)}{3 + 2D(y) \left(\frac{3}{2} + \frac{\chi}{kT}\right)^2}
$$

$$
\frac{\Gamma_2}{\Gamma_2 - 1} = \frac{1}{\nabla_{\text{ad}}} = \left(\frac{\partial \ln P}{\partial \ln T}\right)_{\text{ad}}
$$

Gas heats up

- \rightarrow rapid increase of ionization
- \rightarrow increase in number of free particles
- \rightarrow pressure goes up faster than \sim T

Pressure increases faster than normal In response to temperature increase

$$
\Gamma_3 - 1 = \left(\frac{\partial \ln T}{\partial \ln \rho}\right)_{\text{ad}}
$$

Gas is compressed

- \rightarrow increase in temperature
- \rightarrow rapid increase in ionization
- \rightarrow energy goes into ionizing gas instead of thermal energy

 \rightarrow temperature does not rise as fast

Temperature increases slower than normal in response to compression

Adiabatic exponents can be less than $4/3 \rightarrow$ unstable!

Ionization zones are responsible for stellar pulsations

- second ionization of *He* provides the most significant driving
- ionization zone must be in the radiative region

Cool, luminous stars $(T_{\text{eff}}$ <6300K) will pulsate in an "instability strip" a few hundred K wide on HR diagram

Polytropes

First two structure equations:

$$
\frac{dm}{dr} = 4\pi r^2 \rho
$$

$$
\frac{dP}{dr} = -\rho \frac{Gm}{r^2}
$$

Temperature *T* does not appear in these equations explicitly, but is involved implicitly because pressure *P* usually depends on temperature via the equation of state.

In some cases, however, *P* only depends on density and, not, *T*. in these cases, the above two equations are sufficient to define a solution.

A degenerate gas is such a case.

Polytropes

Assume a "polytropic relation" holds throughout the star:

$$
P = K \rho^{1 + \frac{1}{n}}
$$

- *K* : constant
- *n* : polytropic index

Polytropes are useful in two situations:

- The equation of state is really polytropic Completely degenerate gas
	- Non-relativistic $P \sim \rho^{5/3}$ \rightarrow *n* = 3/2
	- Ultra-relativistic $P \sim \rho^{4/3}$ \rightarrow *n* = 3
- The equation of state + an additional constraint yields a polytropic relation.
	- Isothermal ideal gas $T = T_0$, $P \sim \rho T \sim \rho \implies n = \infty$
	- Fully convective star: convection maintains a fixed *T* gradient

$$
T \sim P^{2/5}
$$
, $P \sim \rho T \sim \rho P^{2/5} \to P \sim \rho^{5/3} \longrightarrow n = 3/2$

Hydrostatic equilibrium

Make this equation dimensionless. First density,

$$
\rho(r) = \rho_c \theta^n(r)
$$
\n
$$
\rho_c = \rho(r=0)
$$
\n
$$
P = K \rho^{1+\frac{1}{n}} \to P = K \rho_c^{1+\frac{1}{n}} \theta^{n+1} = P_c \theta^{n+1} \left(P_c = K \rho_c^{1+\frac{1}{n}}\right)
$$

$$
\frac{1}{r^2}\frac{d}{dr}\left[\frac{r^2}{\rho}\frac{dP}{dr}\right] = -4\pi G\rho \qquad \rho = \rho_c \theta^n \qquad P = P_c \theta^{n+1}
$$

$$
\frac{1}{r^2}\frac{d}{dr}\left[\frac{r^2}{\rho_c\theta^n}\frac{d}{dr}\left(P_c\theta^{n+1}\right)\right] = -4\pi G\rho_c\theta^n
$$
\n
$$
\rightarrow \frac{P_c}{\rho_c}\frac{1}{r^2}\frac{d}{dr}\left[\frac{r^2}{\theta^n}\left(n+1\right)\theta^n\frac{d\theta}{dr}\right] = -4\pi G\rho_c\theta^n
$$
\n
$$
\rightarrow \frac{(n+1)P_c}{4\pi G\rho_c^2}\frac{1}{r^2}\frac{d}{dr}\left[r^2\frac{d\theta}{dr}\right] = -\theta^n
$$

Next, make radius dimensionless:

$$
\xi = \frac{r}{r_n}
$$

$$
r_n^2 \frac{1}{(\xi r_n)^2} \frac{d}{d(\xi r_n)} \left[\left(\xi r_n\right)^2 \frac{d\theta}{d(\xi r_n)} \right] = -\theta^n
$$

$$
\rightarrow \left[\frac{1}{\xi^2} \frac{d}{d\xi} \right] \xi^2 \frac{d\theta}{d\xi} \right] = -\theta^n
$$

Lane-Emden equation

$$
\rightarrow \frac{1}{\xi^2} \left(2\xi \frac{d\theta}{d\xi} + \xi^2 \frac{d^2\theta}{d\xi^2} \right) = -\theta^n
$$

$$
\rightarrow \frac{d^2\theta}{d\xi^2} + \frac{2}{\xi}\frac{d\theta}{d\xi} + \theta^n = 0
$$

$$
\theta'' = -\frac{2}{\xi}\theta' - \theta^n
$$

$$
\theta'' = -\frac{2}{\xi}\theta' - \theta^n \qquad \xi = \frac{r}{r_n} \qquad \boxed{\rho(r) = \rho_c}
$$

$$
\left|\xi = \frac{r}{r_n}\right|
$$

$$
\rho(r)=\rho_c\theta^n(r)
$$

*d*ρ

Boundary conditions

• Center

$$
r = 0 \rightarrow \xi = 0
$$
 $\rho = \rho_c \rightarrow \theta = 1$ $\frac{d\rho}{dr} = 0 \rightarrow \theta' = 0$

 \bullet Surface is at first zero crossing of $\,\theta(\xi)\,$

$$
\rho = 0 \rightarrow \theta = 0 \qquad \xi = \xi_1
$$

Solving Lane-Emden

The Lane-Emden equation can be integrated numerically outward until $\theta = 0$.

- Start at $\xi = 0$. In steps of $d\xi$ compute θ'' , θ' , θ . Stop when $\theta = 0$.
- Get value of ξ_1 , as well as full $\theta(\xi)$ profile.

• Analytic solutions exist for three cases: *n*=0, 1, and 5

$$
n = 0
$$
: $\theta(\xi) = 1 - \frac{\xi^2}{6}$ $\xi_1 = \sqrt{6}$

$$
n = 1: \quad \theta(\xi) = \frac{\sin \xi}{\xi} \qquad \qquad \xi_1 = \pi
$$

$$
n = 5: \quad \theta(\xi) = \left(1 + \frac{\xi^2}{3}\right)^{-1/2} \qquad \qquad \xi_1 = \infty
$$

- Polytropes with *n*>5 have infinite (divergent) mass.
- Only models with *n*=3/2 and *n*=3 are physically relevant.

$$
n = \frac{3}{2}
$$
: $P = K\rho^{5/3}$ $n = 3$: $P = K\rho^{4/3}$

Solving a differential equation numerically

Simplest case: *dy dx* $y = f(x,y)$ Boundary condition: $y(0) = y_0$ $y'(0) = y'_0$

- Choose step Δ*x*
- Start with boundary condition and evaluate y at next step *y*(Δ*x*) = *y*(0) + *y* ′(0)⋅ Δ*x*
- Repeat for all subsequent steps

$$
y(x) = y(x - \Delta x) + \frac{dy}{dx}(x - \Delta x) \cdot \Delta x
$$

Solving a differential equation numerically

• Loop over steps:

$$
y[i] = y[i-1] + \frac{dy}{dx}[i-1] \cdot \Delta x
$$

$$
\frac{dy}{dx}[i] = \frac{dy}{dx}[i-1] + \frac{d^2y}{dx^2}[i-1] \cdot \Delta x
$$

$$
\frac{d^2y}{dx^2}[i] = f(x,y,y')
$$

• As *n* increases, solutions become less centrally concentrated

Polytropes

0

• Total mass enclosed by r ² *dr r* ∫

$$
M(<\xi) = \int_{0}^{r} \rho_c \theta^n 4\pi (\xi r_n)^2 d(\xi r_n) = 4\pi r_n^3 \rho_c \int_{0}^{r} \theta^n \xi^2 d\xi
$$

= $4\pi r_n^3 \rho_c \int_{0}^{r} \xi^2 \left[-\frac{1}{\xi^2} \frac{d}{d\xi} \left(\xi^2 \frac{d\theta}{d\xi} \right) \right] d\xi = 4\pi r_n^3 \rho_c \left[-\xi^2 \frac{d\theta}{d\xi} \right]$

• Total mass in star

$$
M=4\pi r_n^3 \rho_c \left[-\xi_1^2 \theta'(\xi_1)\right]
$$

• Total radius of star

$$
R = r_n \xi_1 = \left[\frac{(n+1)P_c}{4\pi G \rho_c^2} \right]^{1/2} \xi_1
$$

• Pressure - density *Pc*

$$
P_c = K \rho_c^{1+\frac{1}{n}}
$$

Polytropes

We have four equations \rightarrow we can get 4 unknowns.

e.g., if we know the equation of state: *K*, *n*, and the mass *M*, we can

• solve Lane-Emden to get ξ_1 and $\theta'(\xi_1)$

$$
\theta'' = -\frac{2}{\xi}\theta' - \theta^n
$$

• use four equations to get r_n , R , ρ_c , P_c

$$
P_c = K \rho_c^{1+\frac{1}{n}}
$$

$$
R = r_n \xi_1 = \left[\frac{(n+1)P_c}{4\pi G\rho_c^2}\right]^{1/2} \xi_1
$$

$$
M=4\pi r_n^3 \rho_c \left[-\xi_1^2 \theta'(\xi_1)\right]
$$

Polytropes: Structure of a White Dwarf

$$
P_c = K \rho_c^{1+\frac{1}{n}}
$$

$$
M=4\pi r_n^3 \rho_c \left[-\xi_1^2 \theta'(\xi_1)\right]
$$

$$
P_c = K \rho_c^{1+\frac{1}{n}} \qquad M = 4\pi r_n^3 \rho_c \left[-\xi_1^2 \theta'(\xi_1) \right] \qquad R = r_n \xi_1 = \left[\frac{(n+1)P_c}{4\pi G \rho_c^2} \right]^{1/2} \xi_1
$$

Consider a White Dwarf composed of a degenerate, non-relativistic electron gas.

 $\overline{5}$ $\overline{2}$

$$
n=\frac{3}{2},\quad P=K\rho^{5/3}
$$

$$
R^{2} \sim \frac{P_{c}}{\rho_{c}^{2}} \sim \frac{\rho_{c}^{3/3}}{\rho_{c}^{2}} = \rho_{c}^{-1/3} \to \rho_{c} \sim R^{-6}
$$

$$
M \sim r_{n}^{3} \rho_{c} \sim R^{3} R^{-6} \sim R^{-3} \to \frac{R \sim M^{-1/3}}{R \sim M^{-1/3}}
$$

The radius of a White Dwarf shrinks as its mass increases!

Polytropes: Structure of a White Dwarf

$$
R \sim M^{-1/3} \qquad \qquad \rho_c \sim R^{-6}
$$

As the white dwarf mass increases, its radius shrinks and its central density increases.

Eventually, the core will become relativistic.

As the mass keeps growing, the relativistic core grows too.

Polytropes: Structure of a White Dwarf

$$
P_c = K \rho_c^{1+\frac{1}{n}}
$$

$$
M=4\pi r_n^3 \rho_c \left[-\xi_1^2 \theta'(\xi_1)\right]
$$

 $10³$

$$
P_c = K \rho_c^{1+\frac{1}{n}} \qquad M = 4 \pi r_n^3 \rho_c \left[-\xi_1^2 \theta'(\xi_1) \right] \qquad R = r_n \xi_1 = \left[\frac{(n+1) P_c}{4 \pi G \rho_c^2} \right]^{1/2} \xi_1
$$

Consider a White Dwarf composed of a degenerate, ultra-relativistic electron gas. *n* = 3, $P = K \rho^{4/3}$

$$
R^2 \sim \frac{P_c}{\rho_c^2} \qquad \sim \frac{\rho_c^{4/3}}{\rho_c^2} \qquad = \rho_c^{-2/3} \qquad \to \rho_c \sim R^{-3}
$$

$$
M \sim r_n^3 \rho_c \qquad \sim R^3 R^{-3} \qquad \to \qquad \boxed{M = \text{const}}
$$

When the white dwarf is fully relativistic, its mass decouples from its radius and central density!
Polytropes: Structure of a White Dwarf

$$
P_c = K \rho_c^{1+\frac{1}{n}}
$$

$$
M=4\pi r_n^3 \rho_c \left[-\xi_1^2 \theta'(\xi_1)\right]
$$

$$
P_c = K \rho_c^{1+\frac{1}{n}} \qquad M = 4 \pi r_n^3 \rho_c \left[-\xi_1^2 \theta'(\xi_1) \right] \qquad R = r_n \xi_1 = \left[\frac{(n+1) P_c}{4 \pi G \rho_c^2} \right]^{1/2} \xi_1
$$

$$
r_n = \left(\frac{4P_c}{4\pi G\rho_c^2}\right)^{1/2} = \left(\frac{K\rho_c^{4/3}}{\pi G\rho_c^2}\right)^{1/2} = \left(\frac{K}{\pi G}\rho_c^{-2/3}\right)^{1/2} = \left(\frac{K}{\pi G}\right)^{1/2}\rho_c^{-1/3}
$$

$$
M=4\pi r_n^3 \rho_c \left[-\xi_1^2 \theta'\left(\xi_1\right)\right] \quad =4\pi \left(\frac{K}{\pi G}\right)^{3/2} \rho_c^{-1} \rho_c \left[-\xi_1^2 \theta'\left(\xi_1\right)\right]
$$

$$
M = 4\pi \left(\frac{K}{\pi G}\right)^{3/2} \left[-\xi_1^2 \theta'(\xi_1)\right]
$$

Plugging in numbers: $M = 1.46 M_{\odot}$

This is the Chandrasekhar limiting mass for white dwarfs. No white dwarfs are observed with mass greater than this.

Radiative Transfer

Consider energy in the form of light, passing through a medium. The amount of energy passing through depends on location, direction of flow, time, frequency of light, area through which light is passing.

Radiative Transfer

E E_v erg Hz⁻¹ L erg s⁻¹ L_v erg s⁻¹Hz⁻¹ F erg s⁻¹cm⁻² F_{ν} *I*erg s⁻¹cm⁻²st⁻¹ I_v erg s⁻¹cm⁻²st⁻¹Hz⁻¹ $\text{erg s}^{-1}\text{cm}^{-2}\text{Hz}^{-1}$ erg

Energy Specific Energy Luminosity Specific Luminosity Flux Specific Flux **Intensity** Specific Intensity

 $dE = I_v \cdot dt \cdot dA \cdot d\Omega \cdot dV$

Radiative Transfer

 I_{ν} is constant along a ray travelling on a path, whereas flux decreases as *r -2.*

Fewer rays from a given source cross an area dA that is farther away because that area is a smaller fraction of the whole sphere at that radius.

However, an equal number of rays cross the area dA per solid angle because as dA is further away, a fixed solid angle corresponds to a larger area of emitting region by $\sim r^2$.

Radiative Transfer: Emission

Electrons can recombine with atoms or drop to lower energy levels and emit photons

Emission coefficient

$$
\begin{aligned}\nj & \text{erg s}^{-1} \text{cm}^{-2} \text{st}^{-1} \text{cm}^{-1} \\
j_v & \text{erg s}^{-1} \text{cm}^{-2} \text{st}^{-1} \text{Hz}^{-1} \text{cm}^{-1} \\
dE &= j_v \cdot dt \cdot dV \cdot d\Omega \cdot dV\n\end{aligned}
$$

$$
\left| dI_{v} = j_{v} ds \right|
$$

 $j_{\rm\scriptscriptstyle V}$ is the change of $I_{\rm\scriptscriptstyle V}$ per unit length along a path.

We are only considering spontaneous emission here. Stimulated emission depends on *I_v* and is more conveniently treated as a "negative absorption".

Radiative Transfer: Absorption

Absorption

As light passes through a ϵ *ds* medium, it can be absorbed.

$$
dI_{v} = -\alpha_{v} I_{v} ds
$$

 $\alpha_{v}ds$: fractional loss of intensity over path *ds*

 α_{v} : cm⁻¹ positive $\alpha_{v} \rightarrow$ energy loss

The change in intensity depends on intensity itself because the more photons try to move through a medium, the more photons can become absorbed by it.

Radiative Transfer: Absorption

If absorption is due to particles with number density n and cross-sectional area $\sigma_{\rm v}^{\rm v}$

Consider a volume element *ds*⋅ *dA*

Total area covered by particles: $n \cdot dV \cdot \sigma_{v} = n \cdot dA \cdot ds \cdot \sigma_{v}$

Fraction of area that is covered by particles:

$$
\frac{n \cdot dA \cdot ds \cdot \sigma_{v}}{dA} = n \cdot \sigma_{v} \cdot ds = \alpha_{v} ds \rightarrow \boxed{\alpha_{v} = n\sigma_{v}}
$$

Define opacity as:

$$
\boxed{\alpha_{v} = \rho \kappa_{v}}
$$

 $= \rho \kappa_v$ $\kappa_v : \text{ cm}^2 \text{g}^{-1}$

Opacity is a property of the intervening material, like a cross-sectional area per gram of material.

Radiative Transfer: Optical Depth

dI^ν

Define optical depth τ_{v} and $d\tau_{v} = \alpha_{v} ds$

$$
d\tau_{v} = \alpha_{v} ds
$$

$$
dI_{v} = -\alpha_{v} I_{v} ds \longrightarrow \frac{dI_{v}}{ds} = -\alpha_{v} I_{v} \longrightarrow I_{v} = I_{v}(0) e^{-\alpha_{v}s}
$$

$$
dI_{v} = -I_{v} d\tau_{v} \longrightarrow \frac{dI_{v}}{d\tau_{v}} = -I_{v} \longrightarrow I_{v} = I_{v}(0) e^{-\tau_{v}}
$$

Optical depth is the number of e-foldings change in intensity. It is an alternative variable for path.

- Fixed ds = fixed distance
- Fixed $d\tau$ = fixed change of $I_{\rm v}$

Radiative Transfer Equation

Change in intensity over a path length is the sum of emission + absorption (source + sink terms).

$$
dI_{v} = -\alpha_{v}I_{v}ds + j_{v}ds \qquad \rightarrow \qquad \frac{dI_{v}}{ds} = -\alpha_{v}I_{v} + j_{v}
$$

$$
dI_{v} = -I_{v}d\tau_{v} + j_{v}\frac{d\tau_{v}}{\alpha_{v}} \qquad \rightarrow \qquad \frac{dI_{v}}{d\tau_{v}} = -I_{v} + S_{v}
$$

$$
S_{v} = \frac{j_{v}}{\alpha_{v}}
$$
Source function

In stars, the source function is close to the blackbody Planck function.

Radiative Transfer Equation

$$
\frac{dI_{\nu}}{d\tau_{\nu}} + I_{\nu} = S_{\nu} \qquad \rightarrow e^{\tau_{\nu}} \frac{dI_{\nu}}{d\tau_{\nu}} + e^{\tau_{\nu}} I_{\nu} = e^{\tau_{\nu}} S_{\nu}
$$
\n
$$
\rightarrow \frac{d}{d\tau_{\nu}} \Big(I_{\nu} e^{\tau_{\nu}} \Big) = e^{\tau_{\nu}} S_{\nu} \qquad \rightarrow I_{\nu} e^{\tau_{\nu}} = \int_{0}^{\tau_{\nu}} e^{\tau_{\nu}'} S_{\nu} d\tau_{\nu}' + \text{const}
$$

When
$$
\tau_v = 0 \rightarrow I_v = I_v(0) \rightarrow const = I_v(0)
$$

$$
\rightarrow I_{v}e^{\tau_{v}} = I_{v}(0) + \int_{0}^{\tau_{v}} e^{\tau_{v}'} S_{v} d\tau_{v}'
$$

$$
\rightarrow I_{v} = I_{v}(0) e^{-\tau_{v}} + \int_{0}^{\tau_{v}} e^{-(\tau_{v} - \tau_{v}')} S_{v} d\tau_{v}'
$$

Radiative Transfer Equation

$$
I_{v} = I_{v}(0)e^{-\tau_{v}} + \int_{0}^{\tau_{v}} e^{-(\tau_{v}-\tau_{v}')} S_{v} d\tau_{v}'
$$

Final intensity is initial intensity diminished by absorption + integrated source function also diminished by absorption.

For
$$
S_v = \text{const: } I_v(\tau_v) = I_v(0)e^{-\tau_v} + S_v \int_0^{\tau_v} e^{-(\tau_v - \tau_v')} d\tau_v'
$$

= $I_v(0)e^{-\tau_v} + S_v \left[e^{-(\tau_v - \tau_v')} \right]_0^{\tau_v}$

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

$$
\left(1 - e^{-\tau_v}\right) \quad \text{as } \tau_v \to \infty, \quad I_v \to S_v
$$

Radiative Transfer: Net Flux

• If area *dA* is perpendicular to light rays

 $dF_v = I_v d\Omega$

- If normal vector to area dA is at an angle θ to light rays
	- $dF_v = I_v \cos\theta d\Omega$ Flux is reduced because effective area is smaller.
- "Net flux" in the direction of \vec{n} is $\left|F_{_{\rm V}}\right|$

$$
\vec{n} \text{ is } \boxed{F_v(\vec{n}) = \int I_v \cos \theta \, d\Omega}
$$

For an isotropic radiation field intensity is angle-independent.

$$
F_{\rm v}(\vec{n})=I_{\rm v}\int\cos\theta\,d\Omega=0
$$

Within a star there is a net radial flux (outward) causing a departure from isotropy. We can relate this flux to the temperature gradient.

- plane-parallel approximation
- angular dependence through θ only

$$
\bullet \mu = \cos \theta \qquad \qquad \boxed{ds = dz/\mu}
$$

$$
\frac{dI_v}{ds} = -\alpha_v (I_v - S_v) = -\rho \kappa_v (I_v - S_v)
$$

$$
\rightarrow \mu \frac{\partial I_{\nu}}{\partial z} = -\rho \kappa_{\nu} (I_{\nu} - S_{\nu}) \qquad \rightarrow I_{\nu} = S_{\nu} - \frac{\mu}{\rho \kappa_{\nu}} \frac{\partial I_{\nu}}{\partial z}
$$

 $\rho \kappa$ _ν is the fractional loss of I_v per unit length. $(\rho \kappa_{_{\rm \nu}})$ $^{-1}$ is the mean-free-path.

This second term is the change in intensity over the mean-free-path.

This is small $\rightarrow I_{\rm v} \approx S_{\rm v} \approx B_{\rm v}$

To first order:
$$
I_v \approx B_v - \frac{\mu}{\rho K_v} \frac{\partial B_v}{\partial z}
$$

Integrate to get integrate to get $F_v(z) = \int I_v \cos \theta d\Omega = -2π \int I_v(u,z) \mu d\mu$ +1 −1 ∫

$$
d\Omega = 2\pi \sin\theta d\theta = -2\pi d\mu
$$

$$
I_{\nu} \approx B_{\nu} - \frac{\mu}{\rho \kappa_{\nu}} \frac{\partial B_{\nu}}{\partial z} \qquad \qquad F_{\nu}(z) = 2\pi \int_{-1}^{+1} I_{\nu}(\mu, z) \mu d\mu
$$

 B_v is isotropic so only the derivative term has a non-zero angle dependence.

$$
F_{\nu}(z) = 2\pi \int_{-1}^{+1} \left(-\frac{\mu}{\rho \kappa_{\nu}} \frac{\partial B_{\nu}}{\partial z} \right) \mu d\mu = -\frac{2\pi}{\rho \kappa_{\nu}} \frac{\partial B_{\nu}}{\partial z} \int_{-1}^{+1} \mu^{2} d\mu
$$

$$
= -\frac{2\pi}{\rho \kappa_{\nu}} \frac{\partial B_{\nu}}{\partial z} \left[\frac{\mu^{3}}{3} \right]_{-1}^{+1} = -\frac{2\pi}{\rho \kappa_{\nu}} \frac{\partial B_{\nu}}{\partial z} \frac{2}{3}
$$

 $F_{\rm v}(z)$ =

 $3\rho\kappa_{v}$

∂*T*

∂*z*

$$
F_v(z) = -\frac{4\pi}{3\rho\kappa_v} \frac{\partial B_v}{\partial T} \frac{\partial T}{\partial z}
$$

Total flux (integrating over frequency):

$$
F(z) = \int_{0}^{\infty} F_{v}(z) dv = -\frac{4\pi}{3} \frac{\partial T}{\partial z} \int_{0}^{\infty} \frac{1}{\rho \kappa_{v}} \frac{\partial B_{v}}{\partial T} dv
$$

Define the Rosseland mean opacity:

Can be computed for any material, at any density and any temperature (albeit with difficulty).

$$
\frac{1}{\rho \kappa_R} \equiv \frac{\int_{0}^{\infty} \frac{1}{\rho \kappa_v} \frac{\partial B_v}{\partial T} dv}{\int_{0}^{\infty} \frac{\partial B_v}{\partial T} dv}
$$

$$
F(z) = -\frac{4\pi}{3} \frac{\partial T}{\partial z} \int_{0}^{\infty} \frac{1}{\rho \kappa_{v}} \frac{\partial B_{v}}{\partial T} dv
$$

$$
= -\frac{4\pi}{3} \frac{\partial T}{\partial z} \frac{1}{\rho \kappa_R} \int_0^\infty \frac{\partial B_v}{\partial T} dV
$$

$$
= -\frac{4\pi}{3} \frac{\partial T}{\partial z} \frac{1}{\rho \kappa_R} \frac{\partial}{\partial T} \int_0^\infty B_v \, dv
$$

$$
= -\frac{4\pi}{3\rho\kappa_R} \frac{\partial T}{\partial z} \frac{\partial B}{\partial T}
$$

where *B* is the total intensity

Sphere of uniform brightness. At an exterior point,

$$
I = \begin{cases} B, & \theta < \theta_c \\ 0, & \theta > \theta_c \end{cases}
$$

Find flux:
$$
F = \int I \cos\theta d\Omega = B \int_0^{2\pi} d\phi \int_0^{\theta_c} \cos\theta \sin\theta d\theta
$$

\n
$$
= 2\pi B \int_0^{\theta_c} \cos\theta \sin\theta d\theta = 2\pi B \int_0^{\sin\theta_c} x dx = 2\pi B \frac{\sin^2\theta_c}{2}
$$
\n
$$
\sin\theta_c = \frac{R}{r}
$$
\n
$$
F = \pi B \left(\frac{R}{r}\right)^2
$$
\nAt surface: $\boxed{F = \pi B}$

Fabbel We found that the flux coming from an enclosed sphere is: Also, for a blackbody:

$$
F = \pi B
$$

$$
F = \sigma T^4
$$

$$
B = \frac{\sigma T^4}{\pi}
$$

$$
F(z) = -\frac{4\pi}{3\rho\kappa_R} \frac{\partial T}{\partial z} \frac{\partial B}{\partial T}
$$

$$
\frac{\partial B}{\partial T} = \frac{4\sigma T^3}{\pi}
$$

$$
F(z) = -\frac{16\sigma T^3}{3\rho \kappa_R} \frac{\partial T}{\partial z}
$$

Equation of radiative diffusion

• Applies provided that quantities change slowly on scale of mean free path.

• Material properties enter through K_R

Spherical symmetry: $z \rightarrow r$

$$
L_r = F(r) 4\pi r^2
$$

$$
\sigma = \frac{ac}{4}
$$

$$
L_r = -\frac{16\left(\frac{ac}{4}\right)T^3}{3\rho \kappa_R} dr^2
$$

$$
L_r = -\frac{16\pi acr^2T^3}{3\rho\kappa_R}\frac{dT}{dr}
$$

$$
\frac{dT}{dr} = -\frac{3\rho\kappa_{R}L_{r}}{16\pi a c r^{2}T^{3}}
$$

The Eddington Limit

Photons carry momentum \rightarrow absorption of photons must lead to a force.

Spherical symmetric source with luminosity *L*

- *L* 4π*r* $\frac{1}{2}$ (erg s⁻¹cm⁻²) • Energy flux at distance *r* : *L* 4π*cr* • Momentum flux : $\frac{L}{4\pi\sigma r^2}$ (erg cm⁻³) $(E = pc)$
- Multiply by opacity to get force per unit mass

$$
\left(F = \frac{dp}{dt}\right) \qquad F_{\text{rad}} = \frac{\kappa L}{4\pi c r^2} \qquad \left(\text{erg cm g}^{-1}\right)
$$

Opacity is the fraction of momentum flux absorbed per unit mass.

The Eddington Limit

• Inward force per mass due to gravity:

 $F_{\rm grav} =$ *GM r* 2

Radiation balances gravity when $F_{\text{rad}} = F_{\text{grav}}$

$$
\frac{\kappa L}{4\pi c r^2} = \frac{GM}{r^2} \quad \rightarrow \quad L = \frac{4\pi cGM}{\kappa}
$$

At greater luminosities, F_{rad} > F_{grav} and gas will be blown away.

• Assume opacity is due to Thompson scattering by free electrons

$$
\kappa = \frac{\alpha}{\rho} = \frac{n\sigma}{\rho} = \frac{\sigma}{m} = \frac{\sigma_T}{m_H}
$$

$$
L_{\text{edd}} = \frac{4\pi cGMm_H}{\sigma_T} \qquad L_{\text{edd}} = 3.2 \times 10^4 \left(\frac{M}{M_{\odot}}\right) L_{\odot}
$$

The Eddington Limit

$$
L_{\text{edd}} = 3.2 \times 10^4 \left(\frac{M}{M_{\odot}}\right) L_{\odot}
$$

This is the **Eddington limit**

Assumptions:

- Thompson scattering only other opacity sources increase opacity \rightarrow lower L_{edd}
- spherical symmetry
- Mass-Luminosity relation for very massive stars

$$
\left(\frac{L}{L_{\odot}}\right) = 34.2 \left(\frac{M}{M_{\odot}}\right)^{2.4} = L_{\text{edd}} \rightarrow M_{\text{max}} \sim 100 M_{\odot}
$$

Formation of more massive stars cannot be spherically symmetric

Opacity Sources: Electron Scattering

The opacity of a material depends on the composition (*X*,*Y*,*Z*) the temperature T and the density ρ of gas.

$$
\kappa = \kappa_0 \rho^n T^{-s}
$$

1. Electron scattering

$$
\kappa = \frac{n\sigma}{\rho} = \frac{n_e \sigma_e}{\rho}
$$

In an ionized mixture of *H* and *He*: $n_e = \frac{\rho N_A}{\mu_e} = \frac{2}{1+X}$

$$
\kappa_e = \frac{\rho N_A (1+X)}{2} \frac{\sigma_e}{\rho} = \frac{\sigma_e N_A (1+X)}{2}
$$

Opacity Sources: Electron Scattering

If electrons are non-degenerate and non-relativistic, their cross-section is equal to the Thompson cross-section.

$$
\sigma_T = \frac{8\pi}{3} \left(\frac{e^2}{m_e c^2} \right)^2 = 0.6652 \times 10^{-24} \text{ cm}^2
$$

$$
\kappa_e = \frac{\sigma_T N_A (1+X)}{2} \qquad \frac{\kappa_e = 0.2 (1+X) \text{ cm}^2 \text{ g}^{-1}}{}
$$

Cannot use if

- heavy elements are abundant or gas is partially ionized
- \cdot density is high \rightarrow degenerate
- temperature is high \rightarrow relativistic

e opacity has no frequency, density or temperature dependence

$$
\kappa = \kappa_0 \rho^n T^{-s}, \qquad n = s = 0
$$

Opacity Sources: Free-Free Absorption

2. Free-Free Absorption A free electron cannot absorb a photon because energy and momentum cannot both be conserved. However, the presence of a charged ion near the electron can make this possible.

 $\gamma + e^- + i$ on $\rightarrow e^- + i$ on inverse of Bremsstrahlung

$$
\kappa_{ff} \approx 10^{23} \frac{\rho}{\mu_e} \frac{Z_c^2}{\mu_I} T^{-3.5} \text{cm}^2 \text{g}^{-1} \qquad Z_c : \text{ average nuclear charge}
$$

Since free electrons are required, free-free opacity will be negligible for $T < 10^4$ K since *H* will not be ionized.

$$
\kappa_{ff} \approx 4 \times 10^{22} (X+Y)(1+X)\rho T^{-3.5} \text{cm}^2 \text{g}^{-1}
$$

• fully ionized • no metals

$$
\kappa = \kappa_0 \rho^n T^{-s}
$$
, $n = 1$, $s = 3.5$ | Kramers opacity

Opacity Sources: Bound Absorption

3. Bound-Free Absorption

A photon gets absorbed by a bound electron, ionizing it.

$$
\left| \kappa_{\text{bf}} \approx 4 \times 10^{25} Z (1+X) \rho T^{-3.5} \text{cm}^2 \text{g}^{-1} \right|
$$

$$
\bullet
$$
 7>10⁴ K

$$
\kappa = \kappa_0 \rho^n T^{-s}
$$
, $n = 1$, $s = 3.5$ Kramers opacity

4. Bound-Bound Absorption

A photon gets absorbed and causes a transition between bound energy levels in an atom.

- Very complex calculation: absorption line profiles, line broadening.
- ~10 times smaller than f-f or b-f Kramers opacity

Opacity Sources: H- Absorption

5. H Opacity

At low temperature, an extra electron can attach to the H atom.

H has an ionization potential of 0.75 eV so it's very easy to ionize if $T > a$ few thousand K.

$$
\kappa_{H^-} \approx 2.5 \times 10^{-31} \left(\frac{Z}{0.02}\right) \rho^{1/2} T^9 \text{cm}^2 \text{g}^{-1}
$$

$$
3000 < T < 6000 \text{K}
$$

$$
10^{-10} < \rho < 10^{-5} \text{g}
$$

$$
X \sim 0.7, \quad 0.001 < Z < 0.03
$$

Relevant to Sun' s atmosphere!

$$
\kappa = \kappa_0 \rho^n T^{-s}, \qquad n = 0.5, s = -9
$$

In practice, stellar models use opacities calculated using detailed physics.

- As a function of ρ , *T*, *X*, *Y*, *Z* + breakdown of metals
- Calculate all the relevant b-b, b-f, f-f, H⁻, e⁻, scattering effects.
- Get a correct Rosseland mean opacity.
- LANL 1960s following defense calculations
- OPAL Rogers & Iglesias (1992), Iglesias & Rogers (1996)
- Opacity Project Seaton et al. (1994)
- Opacity tables do not cover whole ρ , T , X , Y , Z space. Extrapolation is dangerous.
- Need to do smart interpolation.
- Differences of as much as 30% between projects occur.

Heat Transfer by Conduction

When the density gets very high, degenerate electrons transfer heat by conduction in addition to providing hydrostatic support.

Total energy flux is additive: $F_{\text{tot}} = F_{\text{rad}} + F_{\text{cond}}$

Like resistance in a parallel circuit

- In normal stars, K_{cond} is large \rightarrow conduction is negligible In center of sun: $\kappa_R \sim 0.2$, $\kappa_{\text{cond}} \sim 2 \times 10^9$!!!
- In degenerate dense stars, it can be smaller than $\kappa_{_{R}}$ In center of cool white dwarf: $\kappa_R \sim 0.2$, $\kappa_{\text{cond}} \sim 5 \times 10^{-5}$

Convection

• If luminosity is transported by radiation, then it must obey

$$
L_r = -\frac{16\pi a c r^2 T^3}{3\rho \kappa_R} \frac{dT}{dr}
$$

• In a steady state, the energy transported per time at radius *r* must be equal to the energy generation rate in the stellar interior.

$$
\rightarrow \frac{dT}{dr}
$$
 will be large if $\begin{cases} L \text{ is large} \\ \kappa_R \text{ is large} \end{cases}$

• The temperature gradient cannot be arbitrarily large. If it gets too steep \rightarrow convection takes over as the main mode of energy transport.

Convection: Stability Criteria

- Uniform composition, $T(r)$, $\rho(r)$ profiles
- Displace a mass element by *dr* without exchanging heat with the environment (adiabatically).

- Element expands to maintain pressure balance with its environment.
- Its new density ρ^* and temperature will not in general equal the ambient values at $r + dr$. * and temperature T^*

Convection: Stability Criteria

• Since the change is adiabatic:

$$
\left(\frac{d\ln P}{d\ln \rho}\right)_{\text{ad}} = \Gamma_1 \longrightarrow \frac{\rho}{P} \left(\frac{dP}{d\rho}\right)_{\text{ad}} = \Gamma_1
$$

$$
\rightarrow (d\rho)_{\text{ad}} = \frac{\rho}{\Gamma_1 P} (dP)_{\text{ad}}
$$

$$
\rho^* = \rho(r) + (d\rho)_{\text{ad}} = \rho(r) + \frac{\rho}{\Gamma_1 P} (dP)_{\text{ad}}
$$

$$
= \rho(r) + \frac{\rho}{\Gamma_1 P} \left(\frac{dP}{dr}\right)_{\text{ad}} dr
$$

• Since pressure equilibrium applies:

$$
\left(\frac{dP}{dr}\right)_{\text{ad}} = \frac{dP}{dr}
$$

$$
\rho(r+dr), T(r+dr)
$$
\n
$$
\left(\rho^*, T^*\right) \quad r+dr
$$
\n
$$
\left(\rho(r), T(r)\right)
$$
- If ρ^* > $\rho(r+dr)$: displaced element will be denser than its surroundings and will settle back down \rightarrow STABILITY * > $\rho(r+dr)$
- If $\rho^* < \rho(r + dr)$: buoyancy will cause element to rise further \rightarrow INSTABILITY * $\leq \rho(r+dr)$
- Stability criterion is:

$$
\rho(r) + \frac{\rho}{\Gamma_1 P} \frac{dP}{dr} dr > \rho(r + dr)
$$

$$
\to \frac{\rho(r + dr) - \rho(r)}{dr} < \frac{\rho}{\Gamma_1 P} \frac{dP}{dr}
$$

• Since the change is adiabatic:

$$
\left(\frac{d\ln P}{d\ln T}\right)_{\text{ad}} = \frac{\Gamma_2}{\Gamma_2 - 1} \rightarrow \frac{T}{P} \left(\frac{dP}{dT}\right)_{\text{ad}} = \frac{\Gamma_2}{\Gamma_2 - 1}
$$

$$
\rightarrow (dT)_{\text{ad}} = \frac{\Gamma_2 - 1}{\Gamma_2} \frac{T}{P} (dP)_{\text{ad}}
$$

$$
T^* = T(r) + (dT)_{\text{ad}}
$$

$$
=T(r)+\left(1-\frac{1}{\Gamma_2}\right)\frac{T}{P}(dP)_{\text{ad}}
$$

$$
=T(r)+\left(1-\frac{1}{\Gamma_2}\right)\frac{T}{P}\frac{dP}{dr}dr
$$

- If $T^* < T(r + dr)$: displaced element will be cooler than its surroundings and will settle back down \rightarrow STABILITY
- If T^* > $T(r + dr)$: element will be hotter and thus rise \rightarrow INSTABILITY
- Stability criterion is:

$$
T(r) + \left(1 - \frac{1}{\Gamma_2}\right) \frac{T}{P} \frac{dP}{dr} dr < T(r + dr)
$$

$$
\rightarrow \frac{T(r+dr)-T(r)}{dr} > \left(1-\frac{1}{\Gamma_2}\right)\frac{T}{P}\frac{dP}{dr}
$$

r $r + dr$ $\rho(r)$, $T(r)$ $\rho(r + dr), T(r + dr)$ ρ ∗ $, T^*$

 \cdot Too rapid a change in temperature \rightarrow CONVECTION

$$
\left| \frac{dT}{dr} < \left(1 - \frac{1}{\Gamma_2} \right) \frac{T}{P} \frac{dP}{dr} \right|
$$

• Convert to maximum luminosity that can be carried by radiation

$$
\frac{dT}{dr} = -\frac{3\rho\kappa_R L_r}{16\pi a c r^2 T^3} \qquad \frac{1}{\Gamma_2} \left(1 - \frac{1}{\Gamma_2}\right) \frac{T}{P} \frac{dP}{dr}
$$

$$
\frac{3\rho\kappa_{R}L_{r}}{16\pi a c r^{2}T^{3}} < -\left(1 - \frac{1}{\Gamma_{2}}\right)\frac{T}{P}\frac{dP}{dr}
$$

$$
\frac{dP}{dr} = -\rho \frac{Gm}{r^2}
$$

$$
L_r < \frac{16\pi acG}{3\kappa_R} \left(1 - \frac{1}{\Gamma_2}\right) \frac{T^4 m}{P}
$$

 \cdot If L exceeds this value \rightarrow CONVECTION

- In an unstable region, displaced element is hotter than ambient gas \rightarrow continues to rise
- Eventually, radiation will leak out of the rising gas element.
	- \rightarrow extra energy flux from hotter to cooler regions

• What causes convective instability in some regions of a star?

$$
L_{\max} \sim \frac{1}{\kappa_R} \left(1 - \frac{1}{\Gamma_2} \right) \frac{T^4 m}{P}
$$

When L_{max} is low \rightarrow convection

So, atomic processes + ionization zones \rightarrow convection will happen In regions where *T*~104-105K and *H* is being ionized.

Not in high mass stars

Vulnerable to convection if there is a large luminosity at low mass.

This happens when nuclear energy generation is a very strong function of *T*. CNO in massive stars.

Convection is incredibly difficult to model accurately because fluid motions are very complicated. It is a 3D problem.

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Convection: Energy Flux

- No reliable analytic way to compute
- Lab experiments with fluids are not relevant
	- incompressible
	- too viscous
	- boundaries
- Hydrodynamic simulations are challenging
- Empirical simple model for spherically symmetric stellar models: Mixing Length Theory

1 parameter: length of mixing

Each mass element rises or falls a distance l adiabatically

After one mixing length, The element thermalizes with the local environment.

• *l* should scale with the pressure scale height

$$
\lambda_P = P \left| \frac{dP}{dr} \right|^{-1}
$$
 (length over which pressure changes by 100%)

$$
l = \alpha \cdot \lambda_p
$$
 α : free parameter $(\alpha = 1.6 \pm 0.1)$

 \bullet After rising adiabatically by a distance l , the mass element will be hotter than its surrounding gas by

$$
\Delta T = \left(\left| \frac{dT}{dr} \right| - \left| \frac{dT}{dr} \right|_{\text{ad}} \right) \cdot l \equiv l \cdot \Delta \nabla T
$$

• If the mass element thermalizes at constant pressure, the amount of heat per unit mass released is

$$
\Delta Q = c_p \Delta T = c_p l \Delta \nabla T
$$

• If the average velocity of the mass element is $\overline{\nu}$, the average excess heat flux is

$$
F_{\text{conv}} = \rho \overline{v} \Delta Q = \rho \overline{v} c_p l \Delta \nabla T
$$

$$
\frac{\text{heat}}{\text{mass}} \times \frac{\text{mass}}{\text{volume}} \times \frac{\text{length}}{\text{time}} = \frac{\text{heat}}{\text{area} \cdot \text{time}}
$$

 \bullet The mass element is accelerated due to buoyancy $(\rho < \rho_{\scriptscriptstyle \rm ambient})$

$$
\Delta \rho = \left(\left| \frac{d\rho}{dr} \right| - \left| \frac{d\rho}{dr} \right|_{\text{ad}} \right) l \equiv l \Delta \nabla \rho \quad \rightarrow \overline{\Delta \rho} = \frac{1}{2} l \Delta \nabla \rho
$$

• The average buoyant force per unit volume is

$$
\overline{F} = g \cdot \overline{\Delta \rho} = \frac{1}{2} g l \Delta \nabla \rho
$$
 where $g = \frac{Gm}{r^2}$

• The resulting acceleration is

$$
\overline{a} = \frac{\overline{F}}{\rho} = \frac{gl}{2\rho} \Delta \nabla \rho = \frac{Gml}{2\rho r^2} \Delta \nabla \rho
$$

• If the mass element starts at rest and has a constant acceleration

1

$$
v = \overline{a} \cdot t
$$

\n
$$
t = \frac{l}{\overline{v}} = \frac{l}{\frac{1}{2}v} = \frac{2l}{v}
$$
\n
$$
\rightarrow v = (2\overline{a}l)^{1/2} \rightarrow \overline{v} = \frac{1}{2} (2\overline{a}l)^{1/2}
$$

\n
$$
\rightarrow \overline{v} = \frac{l}{2} \left(\frac{Gm}{\rho r^2} \Delta \nabla \rho \right)^{1/2}
$$

$$
F_{\rm conv} = \rho \overline{\nu} c_P l \Delta \nabla T
$$
 $\qquad \qquad \overline{\nu} =$

$$
\overline{v} = \frac{l}{2} \left(\frac{Gm}{\rho r^2} \Delta \nabla \rho \right)^{1/2}
$$

$$
F_{\text{conv}} = \frac{l}{2} \left(\frac{Gm}{\rho r^2} \Delta \nabla \rho \right)^{1/2} \rho c_P l \Delta \nabla T
$$

$$
\Delta \nabla \rho \equiv \left(\left| \frac{d\rho}{dr} \right| - \left| \frac{d\rho}{dr} \right|_{\text{ad}} \right) \approx \frac{\rho}{T} \left(\left| \frac{dT}{dr} \right| - \left| \frac{dT}{dr} \right|_{\text{ad}} \right) \equiv \frac{\rho}{T} \Delta \nabla T
$$

$$
F_{\text{conv}} = \frac{l}{2} \left(\frac{Gm}{\rho r^2} \frac{\rho}{T} \Delta \nabla T \right)^{1/2} \rho c_p l \Delta \nabla T
$$

$$
F_{\text{conv}} = \frac{l^2}{2} c_P \rho \left(\frac{Gm}{Tr^2}\right)^{1/2} \left(\Delta \nabla T\right)^{3/2}
$$

• How large an excess T gradient is needed to carry luminosity?

$$
\left(\Delta \nabla T\right)^{3/2} = \frac{\frac{L_r}{4\pi r^2}}{\frac{l^2}{2}c_p \rho \left(\frac{Gm}{Tr^2}\right)^{1/2}} = \frac{L_r}{2\pi l^2 c_p \rho r (Gm/T)^{1/2}}
$$

e.g., at *m M* $= 0.5$, $l = \lambda_p = 5 \times 10^9$ cm, solar values for T, ρ, r, L_r, c_p

 $\frac{1}{\sqrt{2}}$

$$
\Delta \nabla T \sim 10^{-10} \text{Kcm}^{-1} \qquad \left| \frac{dT}{dr} \right| \approx \frac{T_c}{R} \sim 10^{-4} \text{Kcm}^{-1}
$$

Requires an excess of only \sim 10⁻⁶ of the temperature gradient!

• When convection occurs in stellar interiors, the resulting temperature gradient equals the adiabatic gradient.

• If composition changes with radius, this may be different. e.g., if μ drops with $r \rightarrow$ heavier material is displaced into lighter material \rightarrow greater stability.

Nuclear burning \rightarrow stabilizing

• When convection is efficient, the exact value of the mixing length *does not matter.*

Near surface, $\lambda_{\stackrel{}{P}}$ is small \rightarrow larger $\Delta \nabla T$ is needed.

• Stellar radius R , surface T , etc. depend on α

x2 change in $\alpha \rightarrow$ few hundred K difference in T_{eff}.

Convective Overshooting Do convective elements overshoot into stable regions due to inertia?

Naively: No because acceleration is small and braking is large in stable zone.

Simulations: Overshooting extends to significant fraction of $\lambda_{\scriptscriptstyle P}^{}$

Alternative to Mixing Length model "Full spectrum of turbulence"

Turbulent processes involve a range of length scales.

It is unclear if this works better than the one-parameter model.

Energy Generation: Gravitational Sources

In a static star, the energy source is thermonuclear

$$
\frac{dL_m}{dm} = \varepsilon
$$

In a contracting/expanding star, if the process is not adiabatic, there is an extra source/sink of energy so that

$$
\frac{dL_m}{dm} \neq \varepsilon \qquad \text{Instead:} \quad \frac{dL_m}{dm} = \varepsilon + \varepsilon_{\text{grav}}
$$

In real stars, contraction/expansion is a local process. e.g., core contracts while envelope expands $\Rightarrow \mathcal{E}_{grav}$ is a function of radius.

$$
\varepsilon_{\text{grav}} = \frac{d\Omega}{dt \cdot dm} = \frac{d}{dt} \left(\frac{Gm}{r} \right) = -\frac{Gm}{r^2} \dot{r}
$$

Energy Generation: Neutrino Losses

 L_m excludes energy flux in neutrinos.

Matter at normal densities and temperatures is completely transparent to neutrinos \rightarrow energy loss

neutrino cross-section: $\sigma_{v} \sim 10^{-44} \varepsilon_{v}^{2}$ cm $(\varepsilon_{v} : \text{energy in MeV})$

$$
\lambda = \frac{1}{n\sigma_v} \qquad \qquad \lambda \sim 10^{20} \varepsilon_v^{-2} \rho^{-1} \text{cm}
$$

It is only possible to get a short mean-free-path if the density is very high. This is only true in the cores of SN where neutrino pressure is important.

$$
\frac{dL_m}{dm} = \varepsilon + \varepsilon_{\text{grav}} - \varepsilon_v
$$

• Lighter nuclei with mass M_i fuse to form a heavier nucleus of mass *My* . The energy liberated is

$$
E = \Delta Mc^2 = \left(\sum_j M_j - M_y\right)c^2
$$

 $4 \times$ ¹H (mass 1.0079 amu) \rightarrow ⁴He (mass 4.0026 amu) $\Delta M \sim 0.7\%$ or original masses, or 26.5MeV

• The mass of a nucleus is not simply equal to the sum of its constituents (i.e., protons and neutrons). There is also binding energy that holds the constituents together.

• The binding energy of a nucleus is defined to be

 $E_B = ($ mass of constituent nucleons – mass of bound nucleus) c^2

$$
E_B = \left[\left(A - Z \right) m_n + Z m_p - M_{\text{nuc}} \right] c^2
$$

 m_n : neutron mass m_p : proton mass

• The binding energy per nucleon is

$$
f = \frac{E_B}{A}
$$

• Binding energy = energy required to separate the nucleus to infinity against binding forces.

• To get energy from fusing lighter nuclei into heavier nuclei, the total binding energy of the light nuclei must be lower than that of the heavier nucleus. $(M_{\text{nuc}}$ for light $> M_{\text{nuc}}$ for heavy)

e.g.,
$$
3 \times {}^{4}He \rightarrow {}^{12}C
$$
 $3 \times {}^{4}He$: $3E_{B,He} = 3 \times (2n + 2p - M_{He})c^{2}$

$$
^{12}C: E_{B,C} = (6n + 6p - M_c)c^2
$$

$$
3M_{He} > M_C \to 3E_{B,He} < E_{B,C} \to \frac{E_{B,He}}{4} < \frac{E_{B,C}}{12}
$$

 E_B for light element *A* • A nuclear reaction can release energy if is less than that for heavy element.

- The most tightly bound nucleus is $56Fe$. Fusion of pure ¹H to 56 Fe yields \sim 8.5MeV per nucleon, the largest part of which (6.6MeV) is already obtained in fusion to 4He.
- Fusion of elements with A<56 yields energy.
- Fission of elements with A>56 yields energy.
- Elements near ⁵⁶Fe are the most tightly bound and are thus not much use for energy production.
- If a star ends up with ⁵⁶ Fe, energy production is over.

Nuclear Energy Generation: Coulomb Barrier

- To fuse, nuclei must surmount a Coulomb barrier.
- Nuclear forces dominate within a radius

 $r_0 = 1.44 \times 10^{-13} A^{1/3} \text{cm}$

• For nuclei of charge Z_1 and Z_2 height of Coulomb barrier is

$$
E_C = \frac{Z_1 Z_2 e^2}{r_0} \sim Z_1 Z_2 \text{ MeV}
$$

• At *T*=10⁷ K, the thermal energy kT is ~10³eV \rightarrow classically there are ZERO particles in a thermal distribution with sufficient energy to fuse at this *T*.

Nuclear Energy Generation: Tunneling

• Quantum tunneling is required for fusion to take place. Tunneling probability is

$$
P = P_0 E^{-1/2} e^{-2\pi \eta} \qquad \eta = \left(\frac{m}{2}\right)^{1/2} \frac{Z_1 Z_2 e^2}{\hbar E^{1/2}}
$$

*P*₀ : depends on nuclei *m* : reduced mass

P increases rapidly with energy *P* decreases with $Z_1, Z_2 \rightarrow$ lightest elements can fuse at lowest temperatures.

• Higher energies and temperatures are needed to fuse heavier nuclei \rightarrow well separated phases in which different elements burn during stellar evolution.

$$
H \rightarrow He
$$

\n
$$
T \downarrow He \rightarrow C, O
$$

\n
$$
C, O \rightarrow Na, Ne, Si, P
$$

• Most thermonuclear reactions in stars proceed through an intermediate state called the "compound nucleus"

 α : projectile (proton or α particle)

 $\alpha + X \rightarrow Z^* \rightarrow Y + \beta$

X : target nucleus

Z∗ : compound nucleus in excited state

- Reactions happen in several steps:
	- Tunneling through the Coulomb barrier
	- Formation of a compound excited nucleus Z*, whose energy depends on the reaction and kinetic energy of reacting particles.
	- Decay of the Z^{*} via emission of photons, neutrons, protons, alpha particles, electrons, etc.

e.g.,
$$
p + {}^{11}B \rightarrow {}^{12}C^* \rightarrow {}^{12}C^{**} + \gamma
$$

$$
\rightarrow {}^{11}B + p
$$

$$
\rightarrow {}^{11}C + n
$$

$$
\rightarrow {}^{12}N + e^- + \overline{v}_e
$$

$$
\rightarrow {}^{8}Be + {}^{4}He
$$

• The probability of a given output depends on the decay lifetime over the sum of all possible lifetimes.

If τ _{*i*} is the lifetime of a particular output, then the probability of that output is:

$$
P_i = \frac{1/\tau_i}{\sum_i 1/\tau_i} = \frac{\tau}{\tau_i} \quad \text{where} \quad \tau = \left(\sum_i 1/\tau_i\right)^{-1} : \text{total mean life of } {}^{12}C^*
$$

• Through the uncertainty principle, the energy width is $\tau_{_i}=\hbar$

$$
P_i = \frac{\Gamma_i}{\sum_j \Gamma_j} = \frac{\Gamma_i}{\Gamma}
$$

where Γ : total energy width of ¹²C^{*}

• The compound nucleus has **Stationary energy levels** corresponding to excited atomic states that can decay via photon emission.

Quasi-stationary levels, which can decay via particles tunneling back through the Coulomb barrier.

- The lifetimes of quasi-stationary states are shorter and they have broader energy ranges.
- At high energy, the width of states increases so that energy levels overlap \rightarrow continuum of excited levels.
Nuclear Energy Generation: Resonances

- The cross-section for some astrophysically interesting reactions depends critically upon the energy level structure of the compound nucleus. At resonant energies, σ can be boosted by orders of magnitude.
- The maximum cross-section is the geometric cross-section $\pi\lambda^2\propto E^{-1}$
- Also, add dominant exponential tunneling factor

$$
\sigma(E) = S(E) E^{-1} e^{-2\pi\eta}
$$

S : astrophysical cross-section: varies very slowly with *E*

- The energy generation rate depends on the energy released per reaction times the reaction rate.
- Which energies of a particle contribute most to the total reaction rate? High *E* gives higher probability for fusion per reaction, but fewer potential reactions
- Why do nuclear reactions have a high *T* dependence?

• Consider particles of type *j* moving with velocity *v* relative to particles of type *k*. The number densities are n_i and n_k .

• The number of reactions per unit time per volume is:

$$
\tilde{r}_{jk} = n_j \cdot n_k \cdot \sigma \cdot v
$$

• To avoid double counting: *r*

$$
\tilde{r}_{jk} = \frac{1}{1 + \delta_{jk}} n_j n_k \sigma v
$$

• Assume that both species have Maxwell-Boltzmann velocity distributions \rightarrow the relative velocity is also Maxwellian.

$$
E = \frac{1}{2}mv^2
$$
, where *m* is the reduced mass: $m = \frac{m_j m_k}{m_j + m_k}$

• The fraction of all pairs with energy between *E* and *E+dE*:

$$
f(E)dE = \frac{2}{\sqrt{\pi}} \frac{E^{1/2}}{(kT)^{3/2}} e^{-E/kT} dE
$$

• The total reaction rate (per volume, per time) is the mono energetic rate integrated over all energies and weighted by *f(E)*

$$
r_{jk} = \frac{1}{1 + \delta_{jk}} n_j n_k \langle \sigma v \rangle
$$

where σv) = σ 0 ∞ \int $\sigma(E) \cdot v \cdot f(E) dE$

- Replace number densities with mass fractions: $X_i \rho = n_i m_i$
- If each reaction releases an amount of energy Q, then:

$$
\varepsilon_{jk} = \frac{1}{1 + \delta_{jk}} \frac{Q}{m_j m_k} \rho X_j X_k \langle \sigma v \rangle
$$
 (energy) times

• All the temperature dependence is contained in $\langle \sigma v \rangle$

$$
\langle \sigma v \rangle = \int_{0}^{\infty} \sigma(E) \cdot v \cdot f(E) dE
$$

$$
\sigma(E) = S(E)E^{-1}e^{-2\pi\eta}
$$

$$
v = \left(\frac{2E}{m}\right)^{1/2}
$$

$$
f(E)dE = \frac{2}{\sqrt{\pi}}\frac{E^{1/2}}{(kT)^{3/2}}e^{-E/kT}dE
$$

$$
= \int_{0}^{\infty} S(E) \, k^{-1} e^{-2\pi \eta} \left(\frac{2}{m}\right)^{1/2} \, k^{1/2} \, \frac{2}{\sqrt{\pi}} \frac{k^{1/2}}{(kT)^{3/2}} \, e^{-E/kT} \, dE
$$

$$
=\frac{2^{3/2}}{(\pi m)^{1/2}}\frac{1}{(kT)^{3/2}}\int\limits_{0}^{\infty}S(E)e^{-2\pi\eta}e^{-E/kT}dE
$$

$$
\langle \sigma v \rangle = \frac{2^{3/2}}{(\pi m)^{1/2}} \frac{1}{(kT)^{3/2}} \int_{0}^{\infty} S(E) e^{-2\pi \eta} e^{-E/kT} dE
$$

$$
\overline{\eta} = 2\pi \eta E^{1/2} = \pi (2m)^{1/2} \frac{Z_1 Z_2 e^2}{\hbar} \qquad \eta = \left(\frac{m}{2}\right)^{1/2} \frac{Z_1 Z_2 e^2}{\hbar E^{1/2}}
$$

$$
\langle \sigma v \rangle = \frac{2^{3/2}}{(\pi m)^{1/2}} \frac{1}{(kT)^{3/2}} \int_{0}^{\infty} S(E) e^{(-E/kT - \overline{\eta}/E^{1/2})} dE
$$

• Ignore *S(E)*. As *E* goes up, the energy/distribution term (Maxwellian) drops, and the tunneling term increases.

$$
\text{Integrand} = S(E)e^{(-E/kT - \overline{\eta}/E^{1/2})}
$$

- Integrand peaks at E_o , where *d*(Integrand)/*dE* = 0
- Since only a narrow range of *E* contributes to the integral, it is ok to assume $S(E) \approx S_0 = {\rm const}$ (for non-resonant reactions)

$$
\frac{d\left(\text{Integrand}\right)}{dE} = e^{\left(-E/kT - \overline{\eta}/E^{1/2}\right)} \left(-\frac{1}{kT} + \frac{\overline{\eta}}{E} \frac{1}{2} E^{-1/2}\right)
$$

 η *E* 1 2 $E^{-1/2} =$ 1 *kT* • This is zero when: $\frac{1}{E} \frac{1}{2} E^{-1/2} = \frac{1}{LT} \longrightarrow E_0 =$ $\overline{\eta}kT$ 2 $\bigg($ \setminus $\left(\begin{array}{c} \bar{\eta}kT \ \overline{\Omega} \end{array}\right)$ ⎠ ⎟ $2/3$

• Evaluate the integral
$$
\int_{0}^{\infty} e^{(-E/kT - \overline{\eta}/E^{1/2})} dE = \int_{0}^{\infty} e^{f(E)} dE
$$

• Expand $f(E)$ about the maximum at $E=E_0$ using a series expansion truncated at the quadratic term

$$
f(E) = f(E_0) + f'(E_0)(E - E_0) + \frac{1}{2}f''(E_0)(E - E_0)^2 + ...
$$

\n
$$
f(E_0) = -\frac{E_0}{kT} - \frac{\overline{\eta}}{E_0^{1/2}} = -\frac{E_0}{kT} - \frac{2E_0}{kT} = -3\frac{E_0}{kT}
$$

\n
$$
f'(E_0) = -\frac{1}{kT} + \frac{\overline{\eta}}{2E_0^{3/2}} = 0
$$

 $\overline{\eta} =$

 $2E_0^{3/2}$

kT

$$
f''(E_0) = -\frac{\overline{\eta}}{2E_0^3} \frac{3}{2} E_0^{1/2} = -\frac{3\overline{\eta}}{4E_0^{5/2}} = -\frac{3}{2E_0 kT}
$$

$$
f(E) = f(E_0) + \frac{1}{2}f''(E_0)(E - E_0)^2 + ...
$$

\n
$$
= -3\frac{E_0}{kT} + \frac{1}{2}\left(-\frac{3}{2E_0kT}\right)(E - E_0)^2 + ...
$$

\n
$$
= -3\frac{E_0}{kT} - \frac{3}{4E_0kT}E_0^2\left(\frac{E}{E_0} - 1\right)^2 + ...
$$

\n
$$
= -3\frac{E_0}{kT} - \frac{3E_0}{4kT}\left(\frac{E}{E_0} - 1\right)^2 + ...
$$

\n
$$
= -\tau - \frac{1}{4}\tau\left(\frac{E}{E_0} - 1\right)^2 + ...
$$

\n
$$
\int_0^{\infty} e^{f(E)} dE = \int_0^{\infty} \exp\left[-\tau - \frac{1}{4}\tau\left(\frac{E}{E_0} - 1\right)^2\right] dE
$$

 Ω

$$
\int_{0}^{\infty} e^{f(E)} dE = \int_{0}^{\infty} \exp\left[-\tau - \frac{1}{4}\tau \left(\frac{E}{E_{0}} - 1\right)^{2}\right] dE
$$

$$
= e^{-\tau} \int_{0}^{\infty} \exp\left[-\frac{\tau}{4}\left(\frac{E}{E_{0}} - 1\right)^{2}\right] dE
$$

 \lfloor

0

 $\|$

4

⎝ $\overline{}$

$$
x = \left(\frac{E}{E_0} - 1\right)
$$

$$
dx = \frac{dE}{E_0}
$$

⎠ ⎟ $\frac{1}{2}$

dE

 \vert

 $E^{\vphantom{\dagger}}_{0}$

 $\left[\begin{array}{cc} 4(E_0) \end{array} \right]$

$$
\int_{0}^{\infty} e^{(-E/kT - \overline{\eta}/E^{1/2})} dE = \frac{\pi^{1/2}}{3} kT \tau^{1/2} e^{-\tau}
$$

• Plug this into the expression for the cross-section

$$
\langle \sigma \nu \rangle = \frac{2^{3/2}}{(\pi m)^{1/2}} \frac{S_0}{(kT)^{3/2}} \int_0^\infty e^{(-E/kT - \bar{\eta}/E^{1/2})} dE
$$

$$
\langle \sigma v \rangle = \frac{2^{3/2}}{(\pi m)^{1/2}} \frac{1}{(kT)^{3/2}} \frac{\pi^{1/2}}{3} S_0 kT \tau^{1/2} e^{-\tau}
$$

$$
\langle \sigma v \rangle = \frac{2^{2/3}}{3(m)^{1/2}} \frac{1}{(kT)^{1/2}} S_0 \tau^{1/2} e^{-\tau}
$$

$$
\tau \sim \frac{E_0}{kT} \sim \frac{(kT)^{2/3}}{kT} = (kT)^{-1/3}
$$

$$
\rightarrow kT \sim \tau^{-3}
$$

$$
\langle \sigma v \rangle \propto S_0 \tau^2 e^{-\tau}
$$

• Peak energy:
$$
E_0
$$

$$
C_0 = \left(\frac{\overline{\eta}kT}{2}\right)^{2/3}
$$

$$
\overline{\eta} = \pi (2m)^{1/2} \frac{Z_1 Z_2 e^2}{\hbar}
$$

$$
\frac{E_0}{kT} = \left(\frac{\overline{\eta}kT}{2}\right)^{2/3} \frac{1}{kT} = \left(\frac{\pi(2m)^{1/2}Z_1Z_2e^2}{2\hbar}\right)^{2/3} (kT)^{-1/3}
$$

$$
m = \frac{m_j m_k}{m_j + m_k} = \frac{A_j A_k}{A_j + A_k} m_p
$$

$$
\frac{E_0}{kT} = \left(\frac{\pi^2 e^4 m_p}{2\hbar^2}\right)^{1/3} \left(\frac{A_j A_k}{A_j + A_k}\right)^{1/3} (Z_1 Z_2)^{2/3} (kT)^{-1/3}
$$

• Peak energy:
$$
\frac{E_0}{kT} = \left(\frac{\pi^2 e^4 m_p}{2\hbar^2}\right)^{1/3} \left(\frac{A_j A_k}{A_j + A_k}\right)^{1/3} (Z_1 Z_2)^{2/3} (kT)^{-1/3}
$$

$$
\frac{E_0}{kT} \approx 6.6 W^{1/3} \left(\frac{T}{10^7 \text{K}}\right)^{-1/3}
$$

The peak energy increases with $W \rightarrow$ separated phases of nuclear burning

$$
W^{1/3} = 10
$$
\n
$$
W = Z_1^2 Z_2^2 \frac{A_j A_k}{A_j + A_k}
$$
\n
$$
W^{1/3} = 10
$$
\n
$$
W^{1/
$$

$$
\varepsilon_{jk} \sim \langle \sigma v \rangle \sim \tau^2 e^{-\tau} \qquad \qquad \tau \sim T^{-1/3}
$$

• Temperature dependence of energy generation rate

$$
\frac{\varepsilon_{jk} - T^{\nu}}{\varepsilon_{jk} - T^{\nu}} \to \nu = \frac{\partial \ln \varepsilon_{jk}}{\partial \ln T} = \frac{\partial \ln \varepsilon_{jk}}{\partial \ln \tau} \frac{\partial \ln \tau}{\partial \ln T}
$$

$$
= \frac{\partial (2 \ln \tau - \tau)}{\partial \ln \tau} \frac{\partial (-\ln T/3)}{\partial \ln T} = \left(2 - \frac{\partial \tau}{\partial \ln \tau}\right) \left(-\frac{1}{3}\right)
$$

$$
= (2 - \tau) \left(-\frac{1}{3}\right) \to \left[\frac{\tau}{\nu} = \frac{\tau}{3} - \frac{2}{3}\right] = 6.6 W^{1/3} \left(\frac{T}{10^{7} \text{K}}\right)^{-1/3} - \frac{2}{3}
$$

 $v \approx 4$ for lightest elements, rising to $v \approx 40$ for heavier nuclei. Strong *T* dependence \rightarrow stars must be stable to *T* fluctuations.

• We can use reaction rates to compute the time derivatives of mass fractions.

per volume, per time:

- Consider the reaction $A + B \rightarrow C$ The abundances of A , B , and C depend on $r_{A,B}$
- Each reaction yields one *C,* so the rate of change of the number density of $C = r_{A,B}$

$$
\frac{dX_C}{dt} = A_C \frac{X_A X_B}{1 + \delta_{AB}} \frac{\rho}{A_A A_B m_p} \langle \sigma v \rangle = A_C \frac{X_A X_B}{1 + \delta_{AB}} r'_{A,B}
$$

$$
\frac{d^{16}O}{dt} = 16\left(^{4}He^{12}C \cdot r'_{4,12}\right)
$$

⎝ $\overline{}$

dt

$$
\frac{d^{12}C}{dt} = 12\left(^{4}He^{8}Be \cdot r'_{4,8} - ^{4}He^{12}C \cdot r'_{4,12}\right)
$$

e.g.,
\n
$$
{}^{4}He + {}^{4}He \rightarrow {}^{8}Be
$$
\n
$$
{}^{8}Be + {}^{4}He \rightarrow {}^{12}C
$$
\n
$$
{}^{12}C + {}^{4}He \rightarrow {}^{16}O
$$

 \overline{a}

⎠ \mathbf{a}

$$
\frac{d^{8}Be}{dt} = 8\left(\frac{{}^{4}He^{4}He}{2} \cdot r'_{4,4} - {}^{4}He^{8}Be \cdot r'_{4,8}\right)
$$

$$
\frac{d^{4}He}{dt} = 4\left(-2\frac{{}^{4}He^{4}He}{2} \cdot r'_{4,4} - {}^{4}He^{8}Be \cdot r'_{4,8} - {}^{4}He^{12}C \cdot r'_{4,12}\right)
$$

 \bullet In "equilibrium", the mass fractions of 8Be \quad and ^{12}C \quad are conserved.

$$
\frac{d^8Be}{dt} = \frac{d^{12}C}{dt} = 0
$$

$$
{}^{4}He + {}^{4}He → {}^{8}Be
$$

$$
{}^{8}Be + {}^{4}He → {}^{12}C
$$

$$
{}^{12}C + {}^{4}He → {}^{16}O
$$

$$
\frac{{}^{4}He^{4}He}{2} \cdot r'_{4,4} = {}^{4}He^{8}Be \cdot r'_{4,8} \rightarrow {}^{8}Be = \frac{1}{2} {}^{4}He \frac{r'_{4,4}}{r'_{4,8}}
$$

$$
{}^{4}He^{8}Be \cdot r'_{4,8} = {}^{4}He^{12}C \cdot r'_{4,12} \rightarrow {}^{12}C = {}^{8}Be \frac{r'_{4,8}}{r'_{4,12}} = \frac{1}{2} {}^{4}He \frac{r'_{4,4}}{r'_{4,12}}
$$

- Nuclear reactions usually involve several steps.
- The rate is controlled by the rate of the slowest reaction in the chain.
- The total energy release is the sum of energies of the individual steps.

The first step in the p-p chain is: $^{}^{1}H+^1H\rightarrow{}^2H+e^++\nu_{e}$

This reaction is non-resonant.

This reaction is slow because it involves a weak decay *While two protons are flashing past each other, one of them undergoes a weak decay into a neutron at that exact instant.*

To make 4He, we need 4 protons, two of which must be converted Into neutrons via either positron decays or e capture.

$$
p \to n + e^+ + v_e \qquad p + e^- \to n + v_e
$$

The reaction rate is:

$$
r_{pp} = 1.15 \times 10^9 T_9^{-2/3} X^2 \rho^2 \exp\left(-3.38/T_9^{1/3}\right) cm^{-3} s^{-1}
$$

The temperature sensitivity is:

$$
v_{pp} = \frac{11.3}{T_6^{1/3}} - \frac{2}{3}
$$
 In sun: $T_6 \approx 15 \rightarrow v_{pp} \approx 4$

The lifetime of a proton against destruction is:

$$
\tau_p = -\frac{n_p}{dn_p/dt} = \frac{n_p}{2r_{pp}} \approx 6 \times 10^9 \,\text{yr}
$$
 For center of Sun

$$
{}^{1}H + {}^{1}H \rightarrow {}^{2}H + e^{+} + v_{e}
$$
\n
$$
pp1 \quad {}^{2}H + {}^{1}H \rightarrow {}^{3}He + \gamma
$$
\n
$$
{}^{3}He + {}^{3}He \rightarrow {}^{4}He + {}^{1}H + {}^{1}H
$$

 $^3He + {^4He} \rightarrow {^7Be} + \gamma$ $^7Be + e^- \rightarrow ^7Li + v_e$ pp2 First two reactions of pp1 $^{7}Li+^{1}H \rightarrow ^{4}He+^{4}He$

 $^7Be+{}^1H \rightarrow {}^8B+\gamma$ ${}^{8}B \rightarrow {}^{8}Be + e^{+} + V_{e} + \gamma$ pp3 First three reactions of pp2 $^8Be \rightarrow ^4He + ^4He$

$$
{}^{1}H + {}^{1}H \rightarrow {}^{2}H + e^{+} + \nu_e
$$
\n
$$
pp1 \quad {}^{2}H + {}^{1}H \rightarrow {}^{3}He + \gamma
$$
\n
$$
{}^{3}He + {}^{3}He \rightarrow {}^{4}He + {}^{1}H + {}^{1}H
$$

First two reactions of pp1
pp2
$$
{}^{3}He + {}^{4}He \rightarrow {}^{7}Be + \gamma
$$

$$
{}^{7}Be + e^{-} \rightarrow {}^{7}Li + \gamma_e
$$

$$
{}^{7}Li + {}^{1}H \rightarrow {}^{4}He + {}^{4}He
$$

 $^7Be+{}^1H \rightarrow {}^8B+\gamma$ ${}^{8}B \rightarrow {}^{8}Be + e^{+} + V_{e} + \gamma$ pp3 First three reactions of pp2 $^8Be \rightarrow ^4He + ^4He$

pp1 is the most direct route but it involves the collision of two short lived 3He nuclei.

Relative importance of chains thus depends on 3He abundance.

As T increases, equilibrium abundance of 3He decreases.

As T increases, importance of pp2 and pp3 relative to pp1 increases.

pp1:
$$
{}^{3}He + {}^{3}He \rightarrow {}^{4}He + {}^{1}H + {}^{1}H
$$

$$
r_{3,3} \sim ({}^{3}He)^{2} \langle \sigma v \rangle_{3,3}
$$
pp2:
$$
{}^{3}He + {}^{4}He \rightarrow {}^{7}Be + \gamma
$$

$$
r_{3,4} \sim {}^{3}He^{4}He \langle \sigma v \rangle_{3,4}
$$
pp2 will dominate when:
$$
r_{3,4} > r_{3,3} \rightarrow \frac{{}^{4}He}{{}^{3}He} \times \frac{\langle \sigma v \rangle_{3,3}}{\langle \sigma v \rangle_{3,4}} \gg 1
$$
pp2:
$$
{}^{7}Be + e^{-} \rightarrow {}^{7}Li + V_{e}
$$

$$
r_{7,-} \sim {}^{7}Be \times e^{-} \langle \sigma v \rangle_{7,-}
$$
pp3:
$$
{}^{7}Be + {}^{1}H \rightarrow {}^{8}B + \gamma
$$

$$
r_{7,+} \sim {}^{7}Be \times p^{+} \langle \sigma v \rangle_{7,+}
$$
pp3 will dominate when:
$$
r_{7,+} > r_{7,-} \rightarrow \langle \sigma v \rangle_{7,+} > \langle \sigma v \rangle_{7,-}
$$

The energy generation rate is:

$$
\varepsilon_{pp} = 2.4 \times 10^4 \frac{\rho X^2}{T_9^{2/3}} \exp\left(-3.38/T_9^{1/3}\right) \text{erg s}^{-1} \text{g}^{-1}
$$

The CNO cycle

$$
{}^{12}C + {}^{1}H \rightarrow {}^{13}N + \gamma
$$

\n
$$
{}^{13}N \rightarrow {}^{13}C + e^{+} + \nu_e
$$

\n
$$
{}^{13}C + {}^{1}H \rightarrow {}^{14}N + \gamma
$$

\nCNO1
$$
{}^{14}N + {}^{1}H \rightarrow {}^{15}O + \gamma
$$

\n
$$
{}^{15}O \rightarrow {}^{15}N + e^{+} + \nu_e
$$

\n
$$
{}^{15}N + {}^{1}H \rightarrow {}^{12}C + {}^{4}He
$$

12 13 14 15 16 17 18 19 F O N C

$$
{}^{15}N + {}^{1}H \rightarrow {}^{16}O + \gamma
$$

CNO2 ${}^{16}O + {}^{1}H \rightarrow {}^{17}F + \gamma$
 ${}^{17}F \rightarrow {}^{17}O + e^{+} + \gamma_e$
 ${}^{17}O + {}^{1}H \rightarrow {}^{14}N + {}^{4}He$

$$
{}^{17}O + {}^{1}H \rightarrow {}^{18}F + \gamma
$$

\n
$$
{}^{18}F \rightarrow {}^{18}O + e^+ + \gamma
$$

\nCNO3
\n
$$
{}^{18}O + {}^{1}H \rightarrow {}^{19}F + \gamma
$$

\n
$$
{}^{19}F + {}^{1}H \rightarrow {}^{16}O + {}^{4}He
$$

The CNO cycle

$$
{}^{12}C + {}^{1}H \rightarrow {}^{13}N + \gamma
$$

\n
$$
{}^{13}N \rightarrow {}^{13}C + e^{+} + \nu_e
$$

\n
$$
{}^{13}C + {}^{1}H \rightarrow {}^{14}N + \gamma
$$

\nCNO1 ${}^{14}N + {}^{1}H \rightarrow {}^{15}O + \gamma$
\n
$$
{}^{15}O \rightarrow {}^{15}N + e^{+} + \nu_e
$$

\n
$$
{}^{15}N + {}^{1}H \rightarrow {}^{12}C + {}^{4}He
$$

This is the slowest reaction and sets the overall rate.

After long enough time, the most abundant nucleus will be 14N Most CNO \rightarrow ¹⁴N

Timescale to reach equilibrium is long

$$
{}^{15}N + {}^{1}H \rightarrow {}^{16}O + \gamma
$$

\nCNO2 ${}^{16}O + {}^{1}H \rightarrow {}^{17}F + \gamma$
\n ${}^{17}F \rightarrow {}^{17}O + e^{+} + \gamma_e$
\n ${}^{17}O + {}^{1}H \rightarrow {}^{14}N + {}^{4}He$

$$
{}^{17}O + {}^{1}H \rightarrow {}^{18}F + \gamma
$$

\n
$$
{}^{18}F \rightarrow {}^{18}O + e^+ + \gamma_e
$$

\nCNO3
\n
$$
{}^{18}O + {}^{1}H \rightarrow {}^{19}F + \gamma_e
$$

\n
$$
{}^{19}F + {}^{1}H \rightarrow {}^{16}O + {}^{4}He
$$

The CNO cycle

The approximate energy generation rate is:

$$
\varepsilon_{CNO} \approx 4.4 \times 10^{25} \frac{\rho X Z}{T_9^{2/3}} \exp\left(-15.228/T_9^{1/3}\right) \text{ergs}^{-1}\text{g}^{-1}
$$

The temperature sensitivity is:

$$
v_{CNO} = \frac{50.8}{T_6^{1/3}} - \frac{2}{3}
$$
 $T_6 \approx 20 \rightarrow v_{CNO} \approx 18$

pp chain vs. CNO cycle

CNO is favored over pp when:

- T is high *Since heavier nuclei are involved*
- Metallicity is high *Since CNO nuclei are needed*

He Burning (triple α)

$$
^{4}He+^{4}He\rightarrow ^{8}Be
$$

This is the inverse of the final reaction in the pp3 chain.

The reaction is endothermic, absorbing 92 keV of energy. Requires about this much energy in Gamow peak

 $\frac{E_0}{kT} \approx 6.6 W^{1/3} T_7^{-1/3}$ *W* = $Z_j^2 Z_k^2$ ² *AjAk* $A_j + A_k$

 $E_0 = 92 \text{keV} \rightarrow T = 1.15 \times 10^8 \text{K}$

Lower limit for He burning

• When $T>10^8$ K, $8Be$ is created roughly as fast as it is destroyed (via inverse reaction)

We can find equilibrium abundance via $\langle \sigma v \rangle$, but we can also use a Saha equation for nuclei:

$$
\frac{n_{\alpha}^2}{n_{s_{Be}}} = \left(\frac{\pi m_{\alpha} kT}{h^2}\right)^{3/2} e^{-Q/kT}
$$
 $Q = 92keV$ $T = 10^8 K \rightarrow \frac{n_{Be}}{n_{\alpha}} = 7 \times 10^{-9}$

$$
{}^{8}Be+{}^{4}He\rightarrow {}^{12}C+\gamma+\gamma
$$

The reaction is exothermic and resonant, proceeding via an excited state ¹²C^{*}

Most ${}^{12}C^*$ decay straight back to ${}^{8}Be + {}^{4}He$, but some emit a photon to form a stable ${}^{12}C$

He Burning

The approximate energy generation rate is:

$$
\varepsilon_{3\alpha} = 5 \times 10^8 \frac{\rho^2 Y^3}{T_9^3} \exp(-4.4/T_9) \text{erg s}^{-1} \text{g}^{-1}
$$

The temperature sensitivity is:

$$
v_{3\alpha} = \frac{4.4}{T_9} - 3 \qquad T_9 = 0.1 \rightarrow v_{3\alpha} \approx 40
$$

He Burning

The effect of He burning on the structure of the star depends on the central conditions at the moment of ignition.

• Low mass stars ($M < 0.4 M_{\text{sun}}$) develop a fully degenerate Helium core before the temperature rises to the He ignition threshold. The core cannot contract or heat up further \rightarrow no He burning \rightarrow He white dwarf

• High mass stars (M > $1.5M_{sun}$) ignite He while the density is well below the degeneracy threshold (ρ <<10 6 g cm⁻³) \rightarrow <u>No large structural changes</u>

• Intermediate mass stars ignite He under conditions of partial degeneracy. This allows a runaway reaction: He flash

He Flash

- Under degenerate conditions, pressure is a function of ρ only, not T.
- Upon He ignition, the energy release leads to a T rise. However, this does not lead to a P rise and hence to expansion and consequent reduction in ρ and T, as in a non-degenrate gas.
- Higher T increases the rate of nuclear reactions further \rightarrow runaway

Non-degenerate core: He ignition $\rightarrow T \uparrow \rightarrow P \uparrow \rightarrow e$ xpand $\rightarrow T \uparrow$ stable

Degenerate core: He ignition \rightarrow T \uparrow \rightarrow ϵ \uparrow \rightarrow T \uparrow unstable

- This continues for a few minutes until T is high enough in the core to lift degeneracy.
- During this time the core radiates with a luminosity similar to the whole Milky Way
- This energy is absorbed by the core and is not detected at the surface.
- The core then expands rapidly until a new stable He burning structure is attained.

Heavier Elements

• Once enough ¹²C is formed by the triple- α reaction, further captures of α particles occur simultaneously with the Carbon forming reaction.

$$
{}^{12}C + {}^{4}He \rightarrow {}^{16}O + \gamma
$$

$$
{}^{16}O + {}^{4}He \rightarrow {}^{20}Ne + \gamma
$$
 (Reactions beyond ²⁰Ne are rare.)

• He burning \rightarrow C, O, Ne. If T>10⁹K, further reactions can occur. Since binding energies of heavy nuclei are comparable, a wide range of reactions are possible.

Summary of Results

1. Mass Conservation

dm dr $= 4\pi r^2$ ρ

dr dm = 1 $4\pi r^2$ ρ

eulerian lagrangian
2. Hydrostatic Equilibrium

eulerian lagrangian

Need equation of state: $P = f(\rho, T, X_i)$

3. Energy Generation

dLr dr $= 4\pi r^2$ ρε

dLm dm $=\mathcal{E}$

eulerian lagrangian

Need nuclear physics: $\varepsilon = f(\rho, T, X_i)$

4. Energy Flow: Radiation

eulerian lagrangian

Need opacity:
$$
\kappa = f(\rho, T, X_i)
$$

4. Energy Flow: Convection

$$
\frac{dT}{dr} = -\left(1 - \frac{1}{\Gamma_2}\right) \frac{Gm\rho T}{r^2 P} \frac{dT}{dm} = -\left(1 - \frac{1}{\Gamma_2}\right) \frac{GmT}{4\pi r^4 P}
$$

eulerian lagrangian

Need adiabatic exponent:
$$
\Gamma_2 = f(\rho, T, X_i)
$$

Equation of State (non-degenerate matter)

$$
P = \frac{1}{3} aT^4 + \frac{N_A k}{\mu} \rho T
$$

Mean molecular weight: 1 μ = 1 $\mu_{_i}$ + 1 $\mu_{_{\scriptscriptstyle{e}}}$ = *Xi* A_i + *i* $\sum \frac{X_i}{A} + \sum \frac{Z_i X_i y_i}{A}$ \overrightarrow{i} *A_i* ∑

Ionization fractions y_i

Saha equation :

$$
\frac{n^{\dagger}n_e}{n^0} = \frac{2G^{\dagger} \left(2\pi m_e kT\right)^{3/2}}{h^3} \exp\left(-\frac{\chi}{kT}\right)
$$

 Ω

e.g., Hydrogen only :
$$
\frac{y^2}{1-y} = \frac{1}{N_A \rho} \frac{\left(2\pi m_e kT\right)^{3/2}}{h^3} \exp\left(-\frac{\chi_H}{kT}\right)
$$

Thermodynamics

Adiabatic index :

$$
\frac{\Gamma_2}{\Gamma_2 - 1} = \left(\frac{\partial \ln P}{\partial \ln T}\right)_{ad}
$$

$$
\Gamma_2 = \frac{32 - 24\beta - 3\beta^2}{24 - 18\beta - 3\beta^2} \qquad \beta = \frac{P_g}{P_{tot}}
$$

(assuming neutral or fully ionized gas)

Opacity: approximations

$$
\frac{1}{\kappa} \approx \frac{1}{\kappa_{H^-}} + \frac{1}{\kappa_e + \kappa_{ff} + \kappa_{bf}}
$$

 $\kappa_e = 0.2(1+X)cm^2g^{-1}$

(assuming fully ionized and no metals)

$$
\kappa_{ff} \approx 4 \times 10^{22} (X+Y)(1+X) \rho T^{-3.5} cm^2 g^{-1}
$$

$$
\kappa_{bf} \approx 4 \times 10^{25} Z (1+X) \rho T^{-3.5} cm^2 g^{-1}
$$

$$
\kappa_{H^-} \approx 2.5 \times 10^{-31} \left(Z/0.02 \right) \rho^{1/2} T^9 cm^2 g^{-1}
$$

Opacity: tables

Convection

Convection happens when:

$$
\left|\frac{dT}{dr}\right|_{rad} > \left|\frac{dT}{dr}\right|_{ad}
$$

$$
L_m > \frac{16\pi acG}{3K_R} \left(1 - \frac{1}{\Gamma_2}\right) \frac{T^4 m}{P}
$$

Energy generation

$$
\varepsilon = \varepsilon_{\text{pp}} + \varepsilon_{\text{CNO}} + \varepsilon_{3\alpha}
$$

$$
\varepsilon_{pp} = 2.4 \times 10^4 \frac{\rho X^2}{T_9^{2/3}} \exp\left(-3.38/T_9^{1/3}\right) \text{ergs}^{-1}\text{g}^{-1}
$$

$$
\varepsilon_{CNO} \approx 4.4 \times 10^{25} \frac{\rho X Z}{T_9^{2/3}} \exp\left(-15.228/T_9^{1/3}\right) \text{ergs}^{-1}\text{g}^{-1}
$$

$$
\varepsilon_{3\alpha} = 5 \times 10^8 \frac{\rho^2 Y^3}{T_9^3} \exp(-4.4/T_9) \text{erg s}^{-1} \text{g}^{-1}
$$

Given M and X,Y,Z, solve structure equations

to get r, P, T, L_m as a function of m

Need to know: $\rho(P,T,X,Y,Z)$ $K_R(\rho, T, X, Y, Z)$ $\Gamma_2(\rho, T, X, Y, Z)$ ε (ρ ,*T*,*X*,*Y*,*Z*)

Some of these equations are indeterminant at center.

It is better to start the integration at a very small $m>0$.

Central boundary conditions

At a very small m>0:

$$
r \neq 0 \qquad m = \frac{4}{3}\pi r^3 \rho_c \to r = \left(\frac{3m}{4\pi \rho_c}\right)^{1/3}
$$

$$
L_m \neq 0 \qquad L_m = \varepsilon_c m
$$

Central boundary conditions

We can get boundary conditions for P and T by expanding and demanding that their derivatives are zero at r=0

 $P(r) = P(0) + P$ ^{*} $\left(0\right)(r - 0) +$ 1 2 $P''(0)(r-0)^2$ $= P_c + P''(0)$ *r* 2 2 0 $P(r) = P_c - \frac{2}{2}$ 3 $\pi G \rho_c^2 r^2$ $P'(r) = -\frac{Gm\rho}{r}$ *r* 2 $=-\frac{G\rho}{2}$ *r* 2 4 3 πr^3 $\bm{\rho}_c$ $\big($ ⎝ $\left(\frac{4}{3}\pi r^3\rho_c\right)^{\gamma}$ ⎠ $\Big) = -\frac{4}{3}$ 3 $\pi G \rho_c^2 r$

Central boundary conditions

• Similarly,

$$
T(r) = T(0) + Y'(0)(r - 0) + \frac{1}{2}T''(0)(r - 0)^2
$$

\n
$$
= T_c + T''(0)\frac{r^2}{2}
$$

\n
$$
T'(r) = -\frac{3\kappa_R \rho L_r}{16\pi acr^2T^3}
$$

\n
$$
T(r) = T_c - \frac{K_c \rho_c^2 \varepsilon_c}{8acT_c^3}r^2
$$

\n
$$
T(r) = T_c - \left(1 - \frac{1}{\Gamma_{2,c}}\right) \frac{2\pi G \rho_c^2 T_c}{3P_c}r^2
$$

Surface boundary conditions

• We can do better than T=P=0 at surface.

guess R and L:

$$
L = 4\pi R^2 \sigma T_s^4 \rightarrow T_s = \left(\frac{L}{4\pi R^2 \sigma}\right)^{1/4}
$$

Guess ρ very small, e.g.,. 10⁻⁵ g cm⁻³:

$$
P_s = \frac{1}{3} a T_s^4 + \frac{N_A k}{\mu} \rho_s T_s
$$

Steps in constructing a stellar model

- 1. Compute ρ , κ , ε , Γ ₂ as a function of P, T, X, Y, Z **DENSITY** 1
	- Beware of P dropping below **OPACITY** 3 aT^4
	- Use approximations for crude results
	- Interpolate using tables (use approximations outside table bounds)
	- ENERGY GENERATION
	- Use formulas for pp, CNO, and 3 α **THERMODYNAMICS**
	- Γ ₂ = 5/3 for ideal gas, 4/3 for radiation pressure
	- Γ ₂ departs from these values when
		- Mixture of ideal gas and radiation
		- Ionization zones

Steps in constructing a stellar model

- 2. Use four structure equations to compute *r*, *P*, *T*, *Lm* vs. *m* given starting values for these (boundary conditions).
	- Given values of $m, r, P, T, L_m, \rho, \kappa, \varepsilon, \Gamma_2$ at shell $\it i$ compute 4 derivatives: *dr* $\frac{d}{dm}$, *dP* $\frac{d}{dm}$, *dLm* $\frac{m}{dm}$, *dT dm*
	- Use these derivatives to compute *r*, *P*, *T*, *Lm* at shell $\,i+1$ **e.g.**, $r[i+1] = r[i] +$ *dr dm* $\bigg($ ⎝ $\left(\frac{dr}{dr}\right)$ ⎠ $\lfloor i \rfloor \times dm$
	- Shell size $dm < 10^{-4} M$ near center and surface.

Steps in constructing a stellar model

- 3. Deal with lack of complete boundary conditions at center or surface.
	- At center, have r, L_m , but not P, T
	- At surface, have P, T, but not r, L_m
	- General approach: guess values for missing conditions at one end, run model, and compare boundary conditions at other end.

PROBLEM: small changes in conditions at center can cause large differences at surface \rightarrow difficult to reach convergence.

SOLUTION: Shoot from both center and surface and meet halfway through star.

- Guess values for T_c and P_c and integrate outwards from $m = 0$ to $m = M/2$
- Guess values for R and L and integrate inwards from $m = M$ to $m = M/2$
- Compute discrepancies Δr , ΔP , ΔT , ΔL_m at $m = M/2$
- Work in log space: log*P*, log*T*, log*r*, log*Lm*

Repeat using new trial values:

 $(\log T_c - d \log T_c)$ and $(\log T_c + d \log T_c)$ with $\log P_c$, $\log R$, $\log L$ $(\log P_c - d \log P_c)$ and $(\log P_c + d \log P_c)$ with $\log T_c$, log*R*, log*L* $(\log R - d \log R)$ and $(\log R + d \log R)$ with $\log P_c$, $\log T_c$, $\log L$ $(\log L - d \log L)$ and $(\log L + d \log L)$ with $\log P_c$, $\log T_c$, $\log R$

- Compute new discrepancies in each case. $\Delta \log r$, $\Delta \log P$, $\Delta \log T$, $\Delta \log L_m$
- Get 16 derivatives. e.g.,

$$
\frac{\partial (\Delta \log r)}{\partial \log P_c} = \frac{(\Delta \log r)_{\log P_c + d \log P_c} - (\Delta \log r)_{\log P - d \log P_c}}{2d \log P_c}
$$

• Use these derivatives to calculate improved boundary conditions.

$$
\log P_c' = \log P_c + \delta \log P_c
$$

$$
\log T_c' = \log T_c + \delta \log T_c
$$

$$
\log R' = \log R + \delta \log R
$$

$$
\log L' = \log L + \delta \log L
$$

• General idea:

$$
-\Delta \log r = \delta \log P_c \frac{\partial (\Delta \log r)}{\partial \log P_c} \to \delta \log P_c = -\Delta \log r \left[\frac{\partial (\Delta \log r)}{\partial \log P_c} \right]^{-1}
$$

• More complicated with 4 variables!

- Invert matrix to solve for $\ \delta \log P_c, \ \delta \log T_c, \ \delta \log R, \ \delta \log L$ (actually take smaller steps ~0.1x)
- Iterate until convergence is reached: discrepancies vanish (i.e., drop below a threshold value)

Gravitational collapse of gas

• Assume a gas cloud of mass M and diameter D

• Sound speed for ideal gas is
$$
c_s = \sqrt{\gamma \frac{P}{\rho}} = \sqrt{\gamma \frac{n k T}{\rho}} = \sqrt{\gamma \frac{k T}{m}}
$$

 t_{sound} = *D* c_s $= D \frac{m}{m}$ γkT $\sqrt{}$ ⎝ $\overline{}$ ⎞ ⎠ ⎟ $1/2$ • Time for sound wave to cross the cloud

• Time for free-fall collapse is
$$
t_{ff} = \frac{1}{\sqrt{G\rho}}
$$

• Gravity beats pressure support when
$$
t_{ff} < t_{sound}
$$

Gravitational collapse of gas

• Critical cloud size is then
$$
t_{ff} = t_{sound} \rightarrow \frac{1}{\sqrt{G\rho}} = D\left(\frac{m}{\gamma kT}\right)^{1/2}
$$

This is the Jeans length

$$
\lambda_J = \left(\frac{\gamma kT}{mG\rho}\right)^{1/2}
$$

• Associated Jeans mass is

$$
M_J = \frac{4}{3}\pi \left(\frac{\lambda_J}{2}\right)^3 \rho \rightarrow
$$

$$
M_J = \frac{\pi}{6} \left(\frac{\gamma kT}{mG} \right)^{3/2} \rho^{-1/2}
$$

Gravitational collapse of gas

• For a typical H_2 molecular cloud:

$$
T = 10 - 100 \text{ K}
$$

$$
n = 10^{2} - 10^{6} \text{ cm}^{-3}
$$

$$
m = 3.34 \times 10^{-24} \text{ g}
$$

$$
\gamma \sim 1
$$

• Associated Jeans mass is

$$
M_J = 70 M_{\odot} \left(\frac{n}{10^3 \text{cm}^{-3}}\right)^{-1/2} \left(\frac{T}{10 \text{K}}\right)^{3/2}
$$

The Orion Nebula

Stars are born in giant gas clouds.

• If the cloud is too hot and not dense enough, it will never collapse. Pressure wins!

• If the cloud is cool and dense enough, it will collapse. Gravity wins!

The Stellar Womb

Stars are born deep in dark molecular clouds:

- cold (10 30 K), dense nebulae
- so cold that molecules (H₂ instead of atomic H) can exist
- dark because visible light cannot penetrate

Stellar Gestation

- something happens to perturb part of a molecular cloud and make it begin to fragment
- as a core of gas collapses, it wants to heat up
- radiates away the excess heat and thus remains cool

Eagle Nebula Pillars
Giant Molecular Gas Cloud

Gravity grows stronger

Gas starts to heat up

Denser \rightarrow Gravity grows stronger

Stellar Gestation

- gets smaller, denser, but not much hotter
- eventually, gas becomes opaque and light escapes less quickly \rightarrow heats up and collapse slows down
- As it heats up, the emitted light moves toward the visible

HST · WFPC2

Star-Birth Clouds · M16 PRC95-44b · ST Scl OPO · November 2, 1995 J. Hester and P. Scowen (AZ State Univ.), NASA

Stellar Gestation

- bursts into view as a visible **protostar**
- hotter, denser, higher pressure
- but still contracting because gravity is stronger too

McNeil'*s nebula*

Denser \rightarrow Gravity grows stronger

Hotter \rightarrow Pressure grows stronger

Denser \rightarrow Gravity grows stronger

Hotter \rightarrow Pressure grows stronger

Stellar Gestation

- The protostar keeps on shrinking until internal pressure can resist gravity
- The protostar collapses until its core reaches 107 K in temperature and fusion starts.
- **Fusion restores hydrostatic** equilibrium. *Hubble*'*s nebula*

The Role of Mass

O stars are most massive $(20-100 M_{sun})$

- Enormous self-gravity, enormous compressive force need enormous pressure to resist gravity
- 10⁷K core temperature is not enough Continue to compress due to gravity, despite fusion
- Compression \rightarrow higher temperature
- Higher temperature \rightarrow faster rate of fusion (larger number of protons have enough energy to fuse)
- Higher fusion rate \rightarrow more pressure
- Equilibrium is reached at very high fusion rate

The Role of Mass

M stars are least massive $(0.08-0.5 M_{sun})$

- Weakest self-gravity, weakest compressive force need less pressure to resist gravity
- Pressure can balance gravity at lower temperature
- Lower temperature \rightarrow lower rate of fusion
- Lower fusion rate \rightarrow lower luminosity

This is the origin of the Mass-Luminosity relation for Main Sequence stars.

$$
L = M^{3.5}
$$
 (in solar units)

Stages of Star Formation on the H-R Diagram

Arrival on the Main Sequence

• The mass of the protostar determines:

- how long the protostellar phase will last
- where the new-born star will land on the MS
- i.e., what spectral type the star will have while on the main sequence

Missing the Main Sequence: Brown Dwarfs

- If the protostar has a mass ≤ 0.08 M_o:
	- It does not contain enough gravitational strength to reach a core temperature of 10^7 K
	- No proton-proton chain fusion reactions occur
	- The object never becomes a star
	- $-$ at 10 $\mathrm{^6}$ K, deuterium fusion begins (but there is not much deuterium) – hydrostatic equilibrium reached
	- this phase is short-lived

The First Brown Dwarf Discovery

Brown Dwarf Gliese 229B

Palomar Observatory Discovery Image October 27, 1994

Hubble Space Telescope Wide Field Planetary Camera 2 November 17, 1995

PRC95-48 · ST Scl OPO · November 29, 1995 T. Nakajima and S. Kulkarni (CalTech), S. Durrance and D. Golimowski (JHU), NASA

Star Formation

Starting Inputs: Mass: 50 Msun Diameter: 0.375 pc Temperature: 10 K Mean mol. Weight: 2.46 (Jeans mass = 1 Msun) Time evolved = 266K years

Initial density and turbulence spectra.

Computing: SPH code, 3.5 M particles 100K CPU-hours on 64 CPUs (65 days) Resolution: 1-5 AU

Bate, M. R., et al. (2002)

Original cloud Denser cloud

Simulation: Matthew Bate, Exeter Visualisation: Richard West, UKAFF

Star Formation

These simulations show:

- Star formation is a very chaotic and dynamic process
- Stars form so close together that they often interact before growing to full size
- Young stars compete for remaining gas with more massive stars
- About half the objects are kicked out of the cluster before they can grow enough to start fusion : brown dwarfs
- Many of the encounters btw. young stars and brown dwarfs strip the dusty disks off the stars suggesting planetary systems could be rare

Matthew Bate University of Exeter Z=3 Z_{\odot}

134606 yr

Matthew Bate University of Exeter

Stellar Initial Mass Function

Evolution of the Sun **Stages in Evolution:**

Sun's Post-Main Sequence Evolutionary Track

Hayashi track

Deuterium burning

Main Sequence $H \rightarrow He$ in core

Red Giant Branch

He core, $H \rightarrow He$ in shell

Tip of the Red Giant Branch Degenerate He core \rightarrow He flash

Horizontal Branch He \rightarrow C,O in core, H \rightarrow He in shell

Asymptotic Giant Branch C,O core, He \rightarrow C,O and H \rightarrow He in shells

Planetary Nebula Not massive enough to burn C,O Sheds outer layers.

White Dwarf Degenerate C,O

Planetary Nebulae

Post - Main Sequence Evolution

Main Sequence Turn-off

