## ASTR-3415-001: Astrophysics Solutions to Exam 1 Spring 2003

 $1.~(40~{
m points~total})$  Assume a gas has an opacity that is constant at all wavelengths and has the functional form

$$\kappa = A \left(\frac{z}{z_0}\right)^{3/2} ,$$

where  $A=3.45\times 10^{11}~{\rm cm^{-1}}$  is the opacity at the reference depth  $z_{\rm o}=5.26\times 10^3~{\rm km}$  and z increases into the gas as viewed from outside. (Note that z increases as  $\tau$  increases.)

(a) [10 points] What is the optical depth of the gas at  $z_{\circ}$ ?

**Solution (a):** Note that in Eq. (I-18) in your notes,

$$d\tau_{\nu} = -\kappa_{\nu} ds$$
,

where  $\tau_{\nu}$  increases in the opposite direction to s. However in this problem, you are told that  $\tau_{\nu}$  and z increase in the same direction (i.e., inward), as such

$$d\tau_{\nu} = \kappa_{\nu} dz$$
.

Now, since  $\kappa$  is independent of wavelength (hence frequency), so is the optical depth. As such,

$$d\tau = \kappa \, dz = A \left(\frac{z}{z_{\circ}}\right)^{3/2} \, dz \ .$$

To determine the optical depth at  $z_{\circ}$ , we just integrate both sides of the equation from the top of the gas, where both  $\tau = 0$  and z = 0, and integrate to  $z_{\circ}$  and solve for  $\tau$  at that position. Then

$$\int_0^{\tau} d\tau = \int_0^{z_\circ} A \left(\frac{z}{z_\circ}\right)^{3/2} dz$$

$$\tau - 0 = \frac{A}{z_\circ^{3/2}} \int_0^{z_\circ} z^{3/2} dz$$

$$\tau = \frac{A}{z_\circ^{3/2}} \frac{2}{5} z^{5/2} \Big|_0^{z_\circ} = \frac{2A}{5} \frac{z_\circ^{5/2}}{z_\circ^{3/2}} = \frac{2A}{5} z_\circ$$

We need to convert  $z_{\circ}$  from km to cm, so  $z_{\circ} = 5.26 \times 10^3$  km  $\times 10^5$  cm/km =  $5.26 \times 10^8$  cm. Using this and the value for A (which is the opacity at  $z_{\circ}$ ), we get

$$\tau = 0.4 (3.45 \times 10^{11} \text{ cm}^{-1}) (5.26 \times 10^8 \text{ cm}) = \boxed{7.26 \times 10^{19}},$$

which is an enormous optical depth (hence we are really in the optically thick regime).

(b) [10 points] At what depth (both in cm and in terms of  $z_{\circ}$ ) does  $\tau=1$ ?

**Solution (b):** Once again, we use the optical depth definition equation but this time we integrate  $\tau$  from 0 to 1 and now solve for z.

$$\int_{0}^{1} d\tau = \int_{0}^{z} A \left(\frac{z}{z_{\circ}}\right)^{3/2} dz$$

$$1 - 0 = \frac{A}{z_{\circ}^{3/2}} \int_{0}^{z} z^{3/2} dz$$

$$1 = \frac{A}{z_{\circ}^{3/2}} \frac{2}{5} z^{5/2} \Big|_{0}^{z} = \frac{2A}{5} \frac{z^{5/2}}{z_{\circ}^{3/2}}$$

$$z^{5/2} = \frac{5z_{\circ}^{3/2}}{2A}$$

$$z = \left(\frac{5z_{\circ}^{3/2}}{2A}\right)^{2/5} = \left(\frac{5(5.26 \times 10^{8} \text{ cm})^{3/2}}{2(3.45 \times 10^{11} \text{ cm}^{-1})}\right)^{2/5}$$

$$z = \left(87.4 \text{ cm}^{5/2}\right)^{2/5} = \boxed{5.98 \text{ cm}}.$$

As such, optical depth unity is reached virtually immediately as we look into the gas — this fog is thick a pea soup!

Now to convert this into units of  $z_{\circ}$ ,

$$z = \frac{5.98 \text{ cm}}{5.26 \times 10^8 \text{ cm}} z_{\circ} = \boxed{1.14 \times 10^{-8} z_{\circ}}.$$

(c) [10 points] Assume the gas is in thermal equilibrium at a temperature of 10,000 K, what is the emissivity (a number is requested here) for the gas at  $z=z_{\circ}$  and  $\lambda=5000$  Å?

**Solution** (c): The emissivity  $\epsilon_{\nu}$  is related to the opacity  $\kappa_{\nu}$  and source function  $S_{\nu}$  by

$$\epsilon_{\nu} = \kappa_{\nu} S_{\nu}$$
.

In thermal equilibrium, the source function is equal to the Planck function, so

$$\epsilon_{\nu} = \kappa_{\nu} B_{\nu} ,$$

as given in Eq. (I-23) in your notes. Since we need frequency and not wavelength here, we must first convert  $\lambda$  to  $\nu$ :

$$\nu = \frac{c}{\lambda} = \frac{2.997925 \times 10^{10} \text{ cm/s}}{5000 \text{ Å} \times 10^{-8} \text{ cm/Å}} = 5.996 \times 10^{14} \text{ Hz}.$$

Now, for a gas at 10,000 K, the Planck function is equal to

$$B_{\nu} = \frac{2h\nu^{3}/c^{2}}{e^{h\nu/k_{B}T} - 1}$$

$$= \frac{2(6.6262 \times 10^{-27} \text{ erg s})(5.996 \times 10^{14} \text{ s}^{-1})^{3}/(2.997925 \times 10^{10} \text{ cm/s})^{2}}{e^{(6.6262 \times 10^{-27} \text{ erg s} \times 5.996 \times 10^{14} \text{ s}^{-1})/(1.3806 \times 10^{-16} \text{ erg/K} \times 10^{4} \text{ K}) - 1}$$

$$= \frac{3.1784 \times 10^{-3} \text{ erg/s/cm}^{2}/\text{Hz/sr}}{e^{2.8777} - 1} = \frac{3.1784 \times 10^{-3} \text{ erg/s/cm}^{2}/\text{Hz/sr}}{16.774}$$

$$= 1.895 \times 10^{-4} \text{ erg/s/cm}^{2}/\text{Hz/sr}.$$

At  $z=z_{\circ}$ , the opacity is

$$\kappa = A \left(\frac{z_{\circ}}{z_{\circ}}\right)^{3/2} = A = 3.45 \times 10^{11} \text{ cm}^{-1},$$

so the emissivity is

$$\epsilon_{\nu} = \kappa_{\nu} B_{\nu} = (3.45 \times 10^{-11} \text{ cm}^{-1})(1.895 \times 10^{-4} \text{ erg/s/cm}^2/\text{Hz/sr})$$
  
=  $6.538 \times 10^7 \text{ erg/s/cm}^3/\text{Hz/sr}$ .

(d) [10 points] What is the value of the source function (again a number is requested) at this temperature and wavelength? Define source function in words.

Solution (d): As shown above, the source function is just the Planck function for gas in thermal equilibrium, hence

$$S_{\nu} = B_{\nu} = 1.895 \times 10^{-4} \text{ erg/s/cm}^2/\text{Hz/sr}$$

I will accept any one of the following two definitions of the source function:

- The ratio of the emissivity to the opacity.
- The ratio of the number of photon creation events to the number of photon destruction events.

- 2. (60 pts) Three terms (see Table I-2 on page I-21 in your notes) exist in an atomic model. Each term is in a subshell that is less than half filled. Term 1 has the following characteristics:  $S=0,\ L=0$  and even parity; Term 2 has  $S=0,\ L=1$ , and odd parity; finally, Term 3 has  $S=1,\ L=2$ , and odd parity. The partition function for the ion of this atom is equal to 1 and the partition function of the atom itself is equal to 4 for a gas at the Sun's effective temperature. The ionization energy of the atom is 8.67 eV.
  - (a) [15 points] Write out the spectroscopic notation for all 5 <u>levels</u> (including J values) arising from these terms.

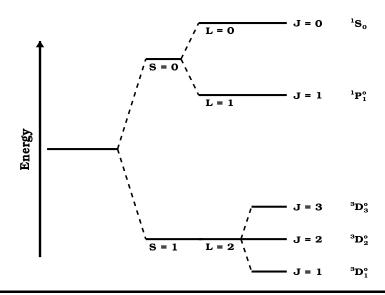
**Solution (a):** As shown in Table I-2 of your notes, a "Term" is a group of levels specified by L and  $S \Longrightarrow$  lines that arise out of the same term are said to belong to the same multiplet. A term will have the number of levels as described by the number of J quantum vectors given by  $|L - S| \le J \le L + S$ . The spectroscopic notation is determined from  ${}^{(2S+1)}L_J$ . We will address each term separately:

- Term 1: Even parity, S=0, (2S+1=1), L=0, (an 'S' state), |L-S|=0, L+S=0, hence J=0.
- Term 2: Odd parity, S=0, (2S+1=1), L=1, (a 'P' state), |L-S|=1, L+S=1, hence J=1.
- Term 3: Odd parity,  $S=1, \ (2S+1=3), \ L=2, \ (a \ ^{\circ}D^{\circ} \ state), \ |L-S|=1, \ L+S=3, \ hence \ J=1,2,3.$  •  $\boxed{^{3}D_{1}^{\circ} \ , \ ^{3}D_{2}^{\circ} \ , \ ^{3}D_{3}^{\circ}}$ .

(b) [15 points] Draw an energy level diagram similar to the one in Figure I-6 on page I-20 of your course notes and locate each level with respect to the other levels using Hund's Rule.

**Solution (b):** We are told from the start that the subshell that each term arises from is less than half filled. As such, Hund's Rules are (1) the higher the S-value, the lower the energy, (2) the higher the L-value (for this S-value), the lower the energy, and (3) the higher the J-value, the higher then energy. The figure on the next page shows the energy diagram that results from these rules for these terms.

## Spectroscopic Notation



(c) [15 points] Determine which of the 7 transitions from these levels are allowed, semi-forbidden, and forbidden. List the reasons for why they are considered as such. (Note that in addition to what is given in your notes, forbidden transitions also occur if either  $\Delta S$  and  $\Delta J$  or  $\Delta L$  and  $\Delta J$  are violated.)

**Solution (c):** The allowed transitions are ones that obey all of the Russell-Saunders selection rules. Violation of these rules will result in either semi-forbidden or forbidden transitions. The table below tallies the results for all the transitions of this model.

Type	Transition	LS Rules Violated
Allowed	$^{1}\mathrm{S}_{0} \rightarrow ^{1}\mathrm{P}_{1}^{\circ}$	all rules obeyed
Semi-forbidden	none	no transitions are semi-forbidden
Forbidden	$^{1}\mathrm{P}_{1}^{\circ} \rightarrow {}^{3}\mathrm{D}_{1}^{\circ}$	no change in parity
	$^{1}\mathrm{P}_{1}^{\circ} \rightarrow {^{3}\mathrm{D}_{2}^{\circ}}$	no change in parity
	$^{1}\mathrm{P}_{1}^{\circ} \rightarrow {}^{3}\mathrm{D}_{3}^{\circ}$	parity, $\Delta S$ and $\Delta J$
	$^{1}\mathrm{S}_{0} \rightarrow {}^{3}\mathrm{D}_{1}^{\circ}$	$\Delta S \ { m and} \ \Delta L$
	$^{1}\mathrm{S}_{0} \rightarrow {}^{3}\mathrm{D}_{2}^{\circ}$	$\Delta S$ and $\Delta L$
	$^{1}\mathrm{S}_{0} \rightarrow {^{3}\mathrm{D}_{3}^{\circ}}$	$\Delta S \ \underline{\text{and}} \ \Delta L$

(d) [15 points] If the ratio of the ion number density to the neutral atom number density is  $4.56\times10^{-2}$  at the continuum formation depth in the Sun's atmosphere, what is the electron density at this depth?

**Solution (d):** This is just an ionization equilibrium question where we use the Saha equation as given in Eq. (I-30) of your notes. Since we have a neutral atom, k = 1, so the Saha equation becomes

$$\frac{N_2}{N_1} n_e = 2 \frac{u_2}{u_1} \frac{(2\pi m k_B T)^{3/2}}{h^3} e^{-\chi_1/k_B T} ,$$

where  $u_1=4$  is the partition function value for the neutral atom (as given in the original question),  $u_2=1$  is the partition function of the ion, and finally,  $\chi_1=8.67~{\rm eV}\times 1.602\times 10^{-12}~{\rm erg/eV}=1.39\times 10^{-11}~{\rm erg}$ . At the continuum formation depth, the Sun's temperature is just its effective temperature (5770 K), so  $k_BT=(1.3806\times 10^{-16}~{\rm erg/K})(5770~{\rm K})=7.966\times 10^{-13}~{\rm erg}$ . Using the following values for our constants, m (the mass of the electron) =  $9.10956\times 10^{-28}~{\rm gm}$ ,  $k_B=1.3806\times 10^{-16}~{\rm erg/K}$ , and  $h=6.6262\times 10^{-27}~{\rm erg}$  s, we can solve the Saha equation for this temperature and ionization energy:

$$\begin{split} \frac{N_2}{N_1} \, n_e &= 2 \frac{1}{4} \, \frac{[2 \pi (9.10956 \times 10^{-28} \text{ gm}) (1.3806 \times 10^{-16} \text{ erg/K}) (5770 \text{ K})]^{3/2}}{(6.6262 \times 10^{-27} \text{ erg s})^3} \, \times \\ &= \frac{e^{-(1.39 \times 10^{-11} \text{ erg})/(7.966 \times 10^{-13} \text{ erg})}}{2.9093 \times 10^{-39} \text{ erg gm})^{3/2}} e^{-17.449} \\ &= 0.5 \, \frac{3.079 \times 10^{-58} \text{ erg}^{3/2} \text{gm}^{3/2}}{2.9093 \times 10^{-79} \text{ erg}^3 \text{s}^3} \times 2.6420 \times 10^{-8} \\ &= 1.3979 \times 10^{13} \text{ erg}^{-3/2} \text{gm}^{3/2} \text{s}^{-3} = 1.3979 \times 10^{13} \text{ (gm cm}^2 \text{ s}^{-2})^{-3/2} \text{gm}^{3/2} \text{s}^{-3} \\ &= 1.3979 \times 10^{13} \text{ cm}^{-3} \, . \end{split}$$

We are told that  $N_2/N_1 = 4.56 \times 10^{-2}$ . Using this fact in the above equation, we can solve for the electron density  $n_e$ :

$$n_e = \frac{1.3979 \times 10^{13} \text{ cm}^{-3}}{N_2/N_1} = \frac{1.3979 \times 10^{13} \text{ cm}^{-3}}{4.56 \times 10^{-2}}$$
$$= 3.066 \times 10^{14} \text{ cm}^{-3}$$
.