## AGNAGN seminar chapter 3. Thermal Equilibrium

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- Refer to original AGNAGN textbook when discussing tables or figures.
- I've tried to fill in every gap I found in original textbook.
- $\bullet\,$  Therefore, this slide consists of 60% original contents and 40% my own interpretation, derivation , or explanation.
- Since 40% my own interpretation may include some incorrect points, I would appreciate it if you tell me any mistake or suspicious point.

• heating : photoionazation (G)

• cooling : recombination(L), cooling by line emission  $\gg$  free-free emission In the equilibrium,

$$G - L = (\text{line}) + (\text{free-free})$$
 (1)

### 3.2 Input by Photoionization

When an ionizing photon  $(h\nu)$  is absorbed, an electron is produced with kinetic energy  $\frac{1}{2}mu^2$ . In this process, energy conservation is

$$h\nu = h\nu_0 + \frac{1}{2}mu^2, \quad h\nu_0 = 13.6 \text{eV}(\text{ionizing energy})$$
 (2)

By integrating this kinetic energy, total energy input by photoionization is

$$G(\mathbf{H}) = \int_{\text{ionizing}} \#(\text{photoelectron}) \times \frac{1}{2} m u^2$$
(3)  
= 
$$\int_{\nu_0}^{\infty} n(\mathbf{H}^0) \#(\text{photon}) h(\nu - \nu_0) a_{\nu}(\mathbf{H}^0) d\nu$$
(4)

where,  $n(\mathrm{H}^0)$ : number density of  $\mathrm{H}^0$ ,  $a_{\nu}(\mathrm{H}^0)$ : cross section. Assume that the  $\mathrm{H}^0$  gas is under the radiation field  $J_{\nu}$ (intensity),

$$\#(\text{photon}) = \frac{4\pi J_{\nu}}{h\nu} \tag{5}$$

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### 3.2 Definition of $T_i$

Therefore, the total energy input (per unit volume, unit time) is

Energy input by photoionization

$$G(\mathbf{H}) = \int_{\nu_0}^{\infty} n(\mathbf{H}^0) \frac{4\pi J_{\nu}}{h\nu} h(\nu - \nu_0) a_{\nu}(\mathbf{H}^0) d\nu$$
(6)

Assume ionization equilibrium (number of  $\mathrm{H}^{0}$  is balanced)

$$n(\mathbf{H}^{0}) \int_{\nu_{0}}^{\infty} \frac{4\pi J_{\nu}}{h\nu} a_{\nu} d\nu = n_{p} n_{e} \alpha_{A}(\mathbf{H}^{0}, T)$$
(7)

the total energy input is written as follows

$$G(\mathbf{H}) = n_e n_p \alpha_A(\mathbf{H}^0, T) \frac{\int_{\nu_0}^{\infty} \frac{4\pi J_\nu}{h\nu} h(\nu - \nu_0) a_\nu(\mathbf{H}^0) d\nu}{\int_{\nu_0}^{\infty} \frac{4\pi J_\nu}{h\nu} a_\nu d\nu}$$
(8)  
=:  $n_e n_p \alpha_A(\mathbf{H}^0, T) \frac{3}{2} k T_i$ (9)

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### 3.2 Interpretation of $T_i$

RHS of eq(9) is the definition of  $T_i$ , which reflects the "feeling" that eq(8) represents the "average" of  $\frac{1}{2}mu^2$  and it corresponds to internal energy. simple example

Consider a simple case;  $J_{\nu} = B_{\nu}(T_{\star}), kT_{\star} < h\nu_0, a_{\nu} \propto \nu^{-3}$ . One can numerically calculate the integrations, and get  $T_i \simeq T_{\star}$ . So, this case is one evidence to regard  $T_i$  as the temperature of photoelectrons.



## 3.2 Hardening( $T_i$ depends on the distance from the star)

We can derive the spatial distribution of  $T_i$ . According to radiative transfer,  $J_{\nu}(r) = J_{\nu}(0)e^{-\tau_{\nu}} = J_{\nu}(0)e^{-na_{\nu}r}$ . Therefore,

$$\frac{3}{2}kT_i(r) = \frac{\int_{\nu_0}^{\infty} \frac{4\pi J_{\nu}(0)}{h\nu} e^{-na_{\nu}r} h(\nu - \nu_0) a_{\nu} d\nu}{\int_{\nu_0}^{\infty} \frac{4\pi J_{\nu}(0)}{h\nu} e^{-na_{\nu}r} a_{\nu} d\nu}$$
(10)

Then we use an inequality (next slide) and we get

$$\frac{3}{2}kT_i(r+dr) = \frac{\int_{\nu_0}^{\infty} \frac{4\pi J_\nu(r)}{h\nu} e^{-na_\nu dr} h(\nu-\nu_0) a_\nu d\nu}{\int_{\nu_0}^{\infty} \frac{4\pi J_\nu(r)}{h\nu} e^{-na_\nu dr} a_\nu d\nu} > \frac{3}{2}kT_i(r)$$
(11)

Note that  $a_{\nu}$  is known to be decreasing (by Fermi's golden rule), so  $e^{-na_{\nu}dr}$  is increasing.

So, at larger distance from the star, mean energy of photoelectrons  $\frac{3}{2}kT_i(r)$  becomes larger.

### 3.2 Lemma (redistribution of probability)

 $\sim$  Lemma(redistribution of probability)  $\cdot$ 

X:random variable

$$P_1(x) = f(x)/Z_1, \quad Z_1 = \int_{\mathbb{R}} f(x)dx$$
 (12)

$$P_2(x) = f(x)g(x)/Z_2, \quad Z_2 = \int_{\mathbb{R}} f(x)g(x)dx$$
 (13)

If g(x) is increasing, then

$$\mathbb{E}_{P_1}[X] < \mathbb{E}_{P_2}[X] \tag{14}$$



Qualitative explanation of this hardening

- $a_{\nu}$ : smaller frequency is effectively absorbed.
- high energy (high frequency) photons penetrate longer distance.
- Therefore, mean energy  $(T_i)$  at larger r becomes higher.

### 3.3 Energy Loss by Recombination

When an electron falls into a quantum state (n, L), its kinetic energy  $\frac{1}{2}mu^2$  is lost. By taking the average of  $\frac{1}{2}mu^2$  in velocity space,

$$\int_0^\infty \frac{1}{2} m u^2 \sigma_{nL}(\mathbf{H}^0, T) u f(u) du \tag{15}$$

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where,  $\sigma_{nL}$ : cross section, f(u): probability density (often Maxwell-Boltzmann distribution is used). Note that  $\sigma_{nL} \cdot u$  means the volume of cylinder, and

$$\sigma_{nL} \cdot u \times f(u) du = \#(\text{recombine with velocity } u)$$
(16)



### 3.3 Energy Loss by Recombination

Suming up mean energies (15) over quantum states (n, L), total energy loss by recombination is

Energy loss by recombination

$$L_R(\mathbf{H}) = n_e n_p k T \beta_A(\mathbf{H}^0, T)$$
(17)

where

$$\beta_A(\mathbf{H}^0, T) = \sum_{n=1}^{\infty} \beta_n(\mathbf{H}^0, T) = \sum_{n=1}^{\infty} \sum_{L=0}^{n-1} \beta_{nL}(\mathbf{H}^0, T)$$
(18)

with

$$\beta_{nL}(\mathbf{H}^{0},T) = \frac{1}{kT} \int_{0}^{\infty} u\sigma_{nL}(\mathbf{H}^{0},T) \frac{1}{2}mu^{2}f(u)du$$
(19)

Note that the denominator kT of eq(19) is just for making it dimensionless.

Since  $\sigma_{nL} \propto u^{-2}$ , electrons with lower kinetic energy are likely to be captured. Therefore,

(mean captured energy) < (mean electron energy)  $\rightarrow \beta_A kT < \frac{3}{2}kT$  (20)

In a pure H nebula that had no radiation losses, the thermal equilibrium is

$$G(\mathbf{H}) = L_R(\mathbf{H}) \tag{21}$$

$$n_e n_p \alpha_A(\mathbf{H}^0, T) \frac{3}{2} k T_i = n_e n_p \beta_A(\mathbf{H}^0, T) k T$$
 (22)

$$\frac{3}{2}kT_i = \beta_A(\mathbf{H}^0, T)kT < \frac{3}{2}kT$$
(23)

Therefore, we get  $T_i < T$ .

In general, the radiation field  $J_{\nu}$  consists of stellar radiation and diffuse radiation  $(J_{\nu} = J_{\nu s} + J_{\nu d})$ . So, we can rewrite  $G(\mathbf{H})$  as follows

$$G(\mathbf{H}) = \int_{\nu_0}^{\infty} n(\mathbf{H}^0) \frac{4\pi (J_{\nu s} + J_{\nu d})}{h\nu} h(\nu - \nu_0) a_{\nu}(\mathbf{H}^0) d\nu =: G_s + G_d$$
(24)

Also, we can decompose  $L_R(\mathbf{H})$  as follows,

$$L_R(\mathbf{H}) = n_e n_p k T(\beta_B + \beta_1), \quad \text{where} \quad \beta_B = \sum_{n=2}^{\infty} \beta_n$$
 (25)

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Note that  $\beta_A = \beta_B + \beta_1$ .

#### on-the-spot approximation

Photon emitted during a recombination to n = 1 level is instantly absorbed by a nearby spot. Energy gain by  $J_{\nu d}$  and energy loss by  $\beta_1$  can simply be omitted.

Using this approximation,

$$G_{\text{OTS}}(\mathbf{H}) = G_s = \int_{\nu_0}^{\infty} n(\mathbf{H}^0) \frac{4\pi J_{\nu s}}{h\nu} h(\nu - \nu_0) a_{\nu}(\mathbf{H}^0) d\nu$$
(26)  
=  $n_e n_p \alpha_B(\mathbf{H}^0, T) \frac{\int_{\nu_0}^{\infty} \frac{4\pi J_{\nu s}}{h\nu} h(\nu - \nu_0) a_{\nu}(\mathbf{H}^0) d\nu}{\int_{\nu_0}^{\infty} \frac{4\pi J_{\nu s}}{\mu\nu} h(\nu - \nu_0) d\nu}$ (27)

 $\int_{\nu_0}^{\infty} \frac{4\pi J_{\nu s}}{h\nu} a_{\nu}(\mathbf{H}^0) d\nu$ 

and

$$L_{\rm OTS}({\rm H}) = n_e n_p k T \beta_B({\rm H}^0, T)$$
(28)

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### 3.3 Rough illustration of on-the-spot approximation



 $E_n = -13.6 \,\mathrm{eV}/n^2$ 

The generalization to include He.

$$G = G(H) + G(He), \quad L_R = L_R(H) + L_R(H_e)$$
 (29)

where

$$G(\text{He}) = n_e n(\text{He}^+) \alpha_A(\text{He}^0, T) \frac{\int_{\nu_2}^{\infty} \frac{4\pi J_{\nu}}{h\nu} h(\nu - \nu_2) a_{\nu}(\text{He}^0) d\nu}{\int_{\nu_2}^{\infty} \frac{4\pi J_{\nu}}{h\nu} a_{\nu}(\text{He}^0) d\nu}$$
(30)

and

$$L_R(\mathbf{H}) = n_e n(\mathbf{H}\mathbf{e}^+) k T \beta_A(\mathbf{H}\mathbf{e}^0, T)$$
(31)

Contributions of other atoms are much less since both G and  $L_R$  are proportional to density n.

The rate of cooling by free-free radiation by charge Z is

$$L_{\rm FF}(Z) = 4\pi j_{\rm ff} \tag{32}$$

$$= \frac{2^5 \pi e^6 Z^2}{3^{3/2} hmc^3} \left(\frac{2\pi kT}{m}\right)^{1/2} g_{\rm ff} n_e n_+ \tag{33}$$

$$= 1.42 \times 10^{-27} Z^2 T^{1/2} g_{\rm ff} n_e n_+ \tag{34}$$

according to  $electromagnetism^1$ .

Here,  $g_{\rm ff}$  is called mean Gaunt factor,  $1.0 < g_{\rm ff} < 1.5$ .

## 3.5 Energy Loss by Collisionally Excited Line Radiation

Ions with energy level gaps  $\sim kT$  such as  $O^+, O^{++}, N^+$  are important sources of cooling.

Here, we consider electrons collide with ions and excite the level 1 to 2.

(energy gap  $\chi = h\nu_{12}$ )

### definition of $\Omega$

Excitation cross section of this process is

$$\sigma_{12}(u) =: \begin{cases} \frac{\pi \hbar^2}{m^2 u^2} \frac{\Omega(1,2)}{\omega_1} & \text{for } \frac{1}{2} m u^2 > \chi \\ 0 & \text{for } \frac{1}{2} m u^2 < \chi \end{cases}$$
(35)

where  $\omega_1$  is the statistical weight of the lower level. Here we define collision strength  $\Omega(1,2)$ .

The fact that  $\sigma \propto u^{-2}$  reflects **Coulomb focusing effect** (quantum scattering theory).

Assume detailed balancing between excitation and deexcitation.

$$n_e n_1 u_1 \sigma_{12}(u_1) f(u_1) du_1 = n_e n_2 u_2 \sigma_{21}(u_2) f(u_2) du_2$$
(36)

Note again that  $\sigma u$  means the volume of the cylinder through which an electron goes.

Also,  $u_1$  and  $u_2$  satisfy energy conservation

$$\frac{1}{2}mu_1^2 = \frac{1}{2}mu_2^2 + \chi \quad (\to u_1 du_1 = u_2 du_2) \tag{37}$$

Then, we set another assumption : Boltzmann distribution

$$\frac{n_2}{n_1} = \frac{\omega_2}{\omega_1} \exp(-\chi/kT) \tag{38}$$

Combining eq(36), (37),(38) and substituting  $f(u) \propto u^2 \exp\left(-\frac{1}{2}mu^2/kT\right)$ ,

## 3.5 Symmetry of $\Omega$

$$\sigma_{12}(u_1)u_1^2 \exp\left(-\frac{1}{2}mu_1^2/kT\right) = \frac{\omega_2}{\omega_1} \exp(-\chi/kT)\sigma_{21}(u_2)u_2^2 \exp\left(-\frac{1}{2}mu_2^2/kT\right)$$
(39)

$$\omega_1 u_1^2 \sigma_{12}(u_1) = \omega_2 u_2^2 \sigma_{21}(u_2) \tag{40}$$

Thus, the deexcitation cross section is derived as

$$\sigma_{21}(u_2) = \frac{\omega_1}{\omega_2} \frac{u_1^2}{u_2^2} \sigma_{12(u_1)} \tag{41}$$
$$\pi \hbar^2 \quad O(1,2)$$

$$=\frac{\pi\hbar^2}{m^2 u_2^2} \frac{\Omega(1,2)}{\omega_2} \tag{42}$$

Remembering the definition of  $\Omega$ , we get

# symmetry of $\Omega$ $\Omega(1,2) = \Omega(2,1) \quad (\Omega \text{ is symmetric}) \qquad (43)$ $\Omega(1,2) = \Omega(2,1) \quad (\Omega \text{ is symmetric}) \quad (43)$

### 3.5 Energy Loss by Collisionally Excited Line Radiation

The total collisional **deexcitation rate** is

$$n_e n_2 q_{21} = n_e n_2 \int_0^\infty u \sigma_{21}(u) f(u) du$$
(44)

$$= n_e n_2 \left(\frac{2\pi}{kT}\right)^{1/2} \frac{\hbar^2}{m^{3/2}} \frac{\Upsilon(1,2)}{\omega_2}$$
(45)

where,  $\Upsilon$  is the velocity-space-averaged collision strength

$$\Upsilon(1,2) = \int_0^\infty \Omega(1,2;E) \cdot \exp(-E/kT) d\left(\frac{E}{kT}\right), \quad \text{with} \quad E = \frac{1}{2}mu_2^2 \qquad (46)$$

Also, we can calculate  $q_{12}$  with ease as

$$q_{12} = \frac{\omega_2}{\omega_1} q_{21} \exp(-\chi/kT)$$
(47)

 $\Upsilon$  must be calculated with **quantum mechanics**.

### 3.5 Relation fine-structure- $\Upsilon$ s satisfy

Now we consider the transition from (S, L, J) to (S', L', J') (fine structure). ; J is coupled angular momentum (satisfies "triangle rule")

$$\Upsilon(SLJ, S'L'J') = (\text{collision strength from } (S, L, J) \text{ to } (S', L', J'))$$
(48)  
$$\Upsilon(SL, S'L') = \sum_{J,J'} \Upsilon(SLJ, S'L'J')$$
(49)



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### 3.5 Relation fine-structure- $\Upsilon$ s satisfy

In general, the summation of  $\Upsilon(SL,S'L')$  is complex. If

- S = 0, then  $|L S| \le J \le L + S \rightarrow J = L$ .
- L = 0, then  $|L S| \le J \le L + S \rightarrow J = S$ .

In those cases, summation is simplified  $\sum_{J,J'} = \sum_{J'}$ .

Assume that all collision strength from  $|S, L, J, m_J\rangle$  to  $|S', L', J', m_{J'}\rangle$  are the same,  $\Upsilon(SLJ, S'L'J') \propto 2J' + 1$  (degeneracy). Therefore, we get

relation between fine-structure- $\Upsilon$ s

$$\Upsilon(SLJ, S'L'J') = \frac{2J'+1}{\sum_{\substack{L+S\\J'=|L-S|}} (2J'+1)} \Upsilon(SL, S'L')$$
(50)  
$$= \frac{2J'+1}{(2S'+1)(2L'+1)} \Upsilon(SL, S'L')$$
(51)

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### example

Consider  ${}^{1}S \rightarrow {}^{3}P$  (Notation;  ${}^{2S+1}L_{J}$ ). The relations are  $\Upsilon({}^{1}S, {}^{3}P_{0}) = \frac{1}{9}\Upsilon({}^{1}S, {}^{3}P)$   $\Upsilon({}^{1}S, {}^{3}P_{1}) = \frac{3}{9}\Upsilon({}^{1}S, {}^{3}P)$ 

$$\Upsilon({}^{1}S, {}^{3}P_{2}) = \frac{5}{9}\Upsilon({}^{1}S, {}^{3}P)$$
(54)

These relations suggest that the rate of the excitation is nearly independent of the distribution of ions among  ${}^{3}P_{0}$ ,  ${}^{3}P_{1}$ , and  ${}^{3}P_{2}$ .

Usage of this relation is rather important in terms of technical reasons.

(52)

(53)

### 3.5 Derivation of cooling rate

We can write down **the detailed balancing** between excitation, deexcitation, and spontaneous emission

$$n_e n_1 q_{12} h \nu_{12} = n_e n_1 q_{21} h \nu_{12} + n_2 A_{21} h \nu_{12}$$
(55)

$$n_e n_1 q_{12} = n_e n_1 q_{21} + n_2 A_{21} \tag{56}$$

Thus,

$$\frac{n_2}{n_1} = \frac{n_e q_{12}}{A_{21}} \left[ 1 + \frac{n_e q_{21}}{A_{21}} \right]^{-1} \tag{57}$$

And therefore

cooling rate

$$L_C = n_2 A_{21} h \nu_{12} = n_e n_1 q_{12} h \nu_{12} \left[ 1 + \frac{n_e q_{21}}{A_{21}} \right]^{-1}$$
(58)

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• 
$$n_e \rightarrow 0$$

$$L_C = n_e n_1 q_{12} h \nu_{12} \left[ 1 + \frac{n_e q_{21}}{A_{21}} \right]^{-1} \to n_e n_1 q_{12} h \nu_{12}$$
(59)

Explaining qualitatively, all collisionally excited electrons are immediately deexcited by spontaneous emission.

•  $n_e \to \infty$ 

$$L_C \to n_e n_1 q_{12} h \nu_{12} \frac{A_{21}}{n_e q_{21}} = \frac{q_{12}}{q_{21}} n_1 A_{21} h \nu_{21}$$

$$= \frac{\omega_2}{\omega_1} e^{-\chi/kT} n_1 A_{21} h \nu_{21} \quad (\text{using eq(47)}) \quad (61)$$

This result is natural considering Boltzmann distribution  $\frac{n_2}{n_1} = \frac{\omega_2}{\omega_1} e^{-\chi/kT}$ .

Some ions (O<sup>++</sup>, N<sup>+</sup>, etc) have more complex energy levels like  ${}^{3}P$ . In such cases, the equilibrium equations are

$$\sum_{j \neq i} n_j n_e q_{ji} + \sum_{j > i} n_j A_{ji} = \sum_{j \neq i} n_i n_e q_{ij} + \sum_{j < i} n_i A_{ij} \quad \text{for } \forall i$$
(62)

And, cooling rate

$$L_C = \sum_i L_C^{(i)} = \sum_i n_i \sum_{j < i} A_{ij} h \nu_{ij}$$

$$\tag{63}$$

n<sub>e</sub> → 0, L<sub>C</sub> becomes a sum of terms like eq(59).
n<sub>e</sub>q<sub>ij</sub> > ∑<sub>k<i</sub> A<sub>ik</sub>, collisional deexcitation is not negligible.

### critical density

$$n_c(i) = \sum_{j < i} A_{ij} / \sum_{j \neq i} q_{ij}$$
(64)

- $n_e < n_c(i)$ , collisional deexcitation of level *i* is negligible.
- $n_e > n_c(i)$ , it is not negligible.

## 3.5 Energy Loss by Collisionally Excited Line Radiation of H

- H<sup>+</sup> has no bound level, no emission.
- H<sup>0</sup> is not so abundant, but may affect the radiative cooling in nebula.
- Important excitation from the ground  $1^2S$ 
  - $2^2 P^0$ . Ly $\alpha$  with  $h\nu = 10.2 \text{eV}$
  - $2^2S$ .  $2\gamma$  decay with  $h\nu' + h\nu'' = 10.2$ eV
- Cross section does not vary as  $u^{-2}$ . It has resonance and peak structure .
- Ys vary fairly slowly.

Table 3.16 Effective collision strengths for H I					
<i>T</i> (K)	1 <sup>2</sup> <i>S</i> , 2 <sup>2</sup> <i>S</i>	$1^{2}S, 2^{2}P^{o}$	1 <sup>2</sup> <i>S</i> , 3 <sup>2</sup> <i>S</i>	$1^2 S, 3^2 P^o$	$1^2S, 3^2D$
10,000	0.29	0.51	0.066	0.12	0.063
15,000	0.32	0.60	0.071	0.13	0.068
20,000	0.35	0.69	0.077	0.14	0.073

Anderson, H., Balance, C. P., Badnell, N. R., & Summers, H. P. 2000, J.Phys.B, 33, 1255.

### 3.5 Resulting Thermal Equilibrium

Including all discussed above, the resulting thermal equilibrium is

$$G = L_R + L_{\rm FF} + L_C \tag{65}$$

• In  $n_e \to 0$ ,  $G, L_R, L_{\rm FF}, L_C$  are all proportional to  $n_e$  and to some ion densities  $n_{\rm ion}$ . So, we can denote  $(G, L_R, L_{\rm FF}, L_C) = n_e n_{\rm ion}(g, l_R, l_{\rm FF}, l_C)$ , where  $(g, l_R, l_{\rm FF}, l_C)$  don't depend on  $n_e$  or  $n_{\rm ion}$ . Then, we get

$$g = l_R + l_{\rm FF} + l_{\rm C} \tag{66}$$

Therefore, resulting T does not depend on  $n_e$  or  $n_{\rm ion}$ .

- In  $n_e > n_c(i)$ , collisional deexcitation is not negligible.  $\rightarrow$  the cooling rate is decreased.
  - $\rightarrow$  the equilibrium temperature is increased.

We set

- $n(O)/n(H) = 7 \times 10^{-4}, n(Ne)/n(H) = 9 \times 10^{-5}, n(N)/n(H) = 9 \times 10^{-5}$
- $\bullet~{\rm O},\,{\rm Ne,N}:80\%$  is singly ionized, 20% is doubly ionized.
- $n({\rm H}^0)/n({\rm H}) = 1 \times 10^{-3}$

The resulting cooling rates are shown in figure 3.2

- dependence of T
  - $kT \ll \chi$ , the contribution is small.
  - $kT \sim \chi$ , it increases rapidly.
  - $kT \gg \chi$ , it decreases slowly.
- low T: O<sup>++</sup> has the greatest contribution.
- high  $T: O^+$  has the greatest contribution.
- The contribution of  $H^0$  is small in all T.

## 3.5 Figure 3.2



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(effective heating rate) = 
$$G - L_R = L_{FF} + L_C$$
 (67)

- In figure 3.2, intersection of dashed  $\operatorname{curve}(G L_R)$  and solid  $\operatorname{curve}(L_C + L_{\mathrm{FF}})$  is the solution of equilibrium, and its typical value is  $T \sim 7000$ K.
- At high  $n_e$ , cooling is suppressed by collisional deexcitation and result in T well above that at low  $n_e$ .

### Appendix: Proof of the lemma

$$\mathbb{E}_{P_{2}}[X] - \mathbb{E}_{P_{1}}[X] = \frac{1}{Z_{2}} \int_{\mathbb{R}} xf(x)g(x)dx - \frac{1}{Z_{1}} \int_{\mathbb{R}} xf(x)dx$$
(68)  
$$= \frac{1}{Z_{1}Z_{2}} \bigg[ Z_{1} \int_{\mathbb{R}} xf(x)g(x)dx - Z_{2} \int_{\mathbb{R}} xf(x)dx \bigg]$$
(69)  
$$= \frac{1}{Z_{1}Z_{2}} \bigg[ \int_{\mathbb{R}^{2}} xf(x)g(x)f(y)dxdy - \int_{\mathbb{R}^{2}} xf(x)g(y)f(y)dxdy \bigg]$$
(70)

Regarding double integration

$$\int_{\mathbf{R}^2} f(x)f(y)[xg(x) - xg(y)]dxdy =: I$$
(71)

we use symmetry trick

$$I = \frac{1}{2} \int_{\mathbb{R}^2} f(x) f(y) [xg(x) - xg(y) + yg(y) - yg(x)] dxdy$$
(72)

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$$I = \frac{1}{2} \int_{\mathbf{R}^2} f(x) f(y) (x - y) (g(x) - g(y)) dx dy$$
(73)

If g(x) is increasing,

$$x - y > 0 \quad \Rightarrow \quad g(x) - g(y) > 0$$

$$\tag{74}$$

and therefore

$$I > 0 \tag{75}$$

$$\mathbb{E}_{P_2}[X] - \mathbb{E}_{P_1}[X] = \frac{I}{Z_1 Z_2} > 0$$

$$\mathbb{E}_{P_2}[X] > \mathbb{E}_{P_1}[X]$$
(76)
(77)

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