

Homework 6 Solutions

Schroeder 3.5, 3.8, 3.9, 3.10, ~~3.11~~
3.14, 3.25 a-e.

Carter 7-4, 7-8, 7-11, 7-13, 7-15

Schroeder:

$$3.5 \quad \Omega = \left(\frac{eN}{q}\right)^q \quad S = k \ln \left(\frac{eN}{q}\right)^q = kq \ln \left(\frac{eN}{q}\right)$$

$$S = kq (\ln e + \ln N - \ln q) = kq (\ln N - \ln q + 1)$$

$U = q \cdot \epsilon$, where ϵ is the size of each energy unit.

$$S = k \frac{U}{\epsilon} (\ln N - \ln U + \ln \epsilon + 1)$$

$$\left(\frac{1}{T}\right) = \frac{\partial S}{\partial U} = \frac{k}{\epsilon} \left[\ln \left(\frac{N\epsilon}{U}\right) + 1 \right] + \frac{kU}{\epsilon} \cdot \frac{-1}{U} = \left(\frac{k}{\epsilon} \ln \left(\frac{N\epsilon}{U}\right) \right)$$

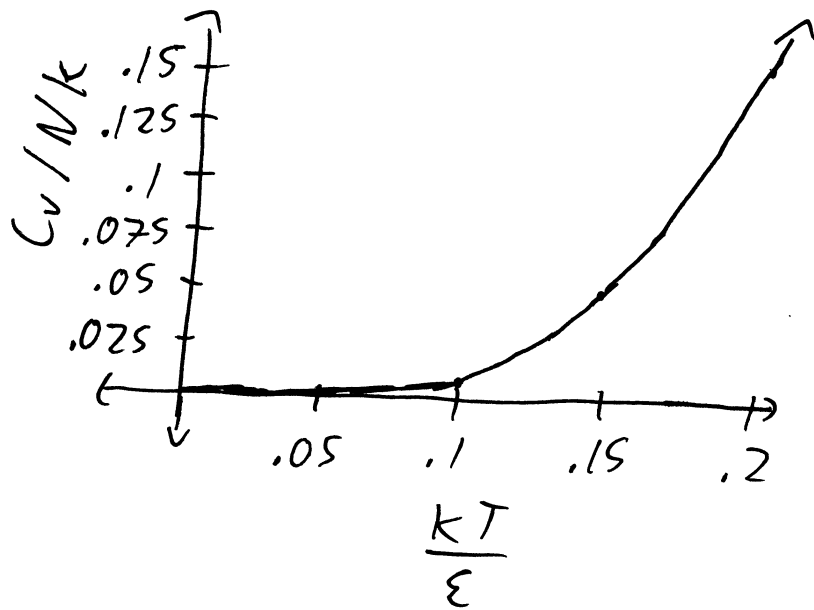
Solving for U yields

$$U = N\epsilon e^{-\epsilon/kT}$$

$U @ T=0 \rightarrow 0$,
which makes sense.

$$3.8 \quad C_v = N \epsilon \frac{d}{dT} e^{-\epsilon/kT}$$

$$= N \epsilon \frac{\epsilon}{kT^2} e^{-\epsilon/kT} = N k \left(\frac{\epsilon}{kT} \right)^2 e^{-\epsilon/kT}$$



... approximately.

3.9 Multiplicity = 2^N (like
2-state magnet)

$$S = k \ln(2^N) = \boxed{Nk \ln 2}$$

$$\text{For } N = 6.02 \cdot 10^{23}, \quad \frac{S}{k} = 4.2 \cdot 10^{23}$$

$$S = 5.8 \text{ J/K}$$

$$3.10 \text{ a. } \Delta S = \frac{Q}{T} = \frac{mL}{T} = \frac{30 \text{ g } 333 \text{ J/g}}{273 \text{ K}} = \boxed{36.6 \text{ J/K}}$$

$$\text{b. } \Delta S = \int_{T_i}^{T_f} \frac{C dT}{T} = C \ln\left(\frac{T_f}{T_i}\right) \\ = (30 \text{ g}) \left(4.186 \frac{\text{J}}{\text{g} \cdot \text{K}}\right) \ln\left(\frac{298}{273}\right) = \boxed{11.0 \frac{\text{J}}{\text{K}}}$$

c. Heat lost by kitchen = heat gained by water!

$$\Delta S = \frac{Q_{\text{kitchen}}}{T_{\text{kitchen}}} = \frac{mL + mc\Delta T}{T_{\text{kitchen}}} \\ = \frac{-(30 \text{ g})(333 \text{ J/g}) - (30 \text{ g}) \cdot 4.186 \frac{\text{J}}{\text{g} \cdot \text{K}} \cdot 25 \text{ K}}{298 \text{ K}} \\ = \boxed{-44.1 \text{ J/K}}$$

$$\text{d. } \Delta S_{\text{universe}} = 36.6 + 11.0 - 44.1 \\ = \boxed{3.5 \text{ J/K}}$$

$$3.14 \quad \Delta S = \int_0^{T_f} \frac{C_v}{T} dT = \int_0^{T_f} \frac{aT + bT^3}{T} dT$$

$$S(T_f) = aT_f + \frac{b}{3} T_f^3 + C$$

$$S(0) = 0 \Rightarrow C = 0$$

$$S(T) = aT + \frac{b}{3} T^3$$

$$S(1K) = (0.00135 \frac{J}{K^2}) \cdot (1K) + \frac{1}{3} (0.0000298 \frac{J}{K^4}) (1K)^3$$

$$= 0.00136 J/K$$

$$S(10K) = 0.0218 J/K$$

$$\frac{S(1K)}{K} = 9.8 \cdot 10^{19}$$

$$\frac{S(10K)}{K} = 1.58 \cdot 10^{21}$$

3.25

a. $S = k \ln \Omega$

$$= k \ln \left(\frac{q+N}{q} \right)^q + k \ln \left(\frac{q+N}{N} \right)^N$$

$$S = kq \ln \left(\frac{q+N}{q} \right) + kN \ln \left(\frac{q+N}{N} \right)$$

(Factors of order $\sqrt{N}$ or $\sqrt{q}$ are considered small compared to factors of order q & N)

b. $\frac{1}{T} = \frac{\partial S}{\partial U} = \frac{\partial q}{\partial U} \frac{\partial S}{\partial q} = \frac{1}{\epsilon} \frac{\partial S}{\partial q}$ (since $q\epsilon = U$)

$$= \frac{k}{\epsilon} \frac{\partial}{\partial q} \left(q \ln(q+N) - q \ln q + N \ln(q+N) - N \ln N \right)$$

$$= \frac{k}{\epsilon} \left(\ln(q+N) + \frac{q}{q+N} - \ln q - \frac{q}{q} + \frac{N}{q+N} + 0 \right)$$

$$= \frac{k}{\epsilon} \left(\ln \left(\frac{q+N}{q} \right) + \frac{q+N}{q+N} - \frac{q}{q} \right)$$

$$\frac{1}{T} = \frac{k}{\epsilon} \ln \left(1 + \frac{N}{q} \right)$$

$$T = \frac{\epsilon}{k \ln\left(1 + \frac{N}{q}\right)} = \frac{\epsilon}{k \ln\left(1 + \frac{N\epsilon}{u}\right)}$$

c. Solving the above equation for u in terms of T yields

$$u = \frac{N\epsilon}{e^{\epsilon/kT} - 1}$$

$$C = \frac{\partial u}{\partial T} = \frac{-N\epsilon}{(e^{\epsilon/kT} - 1)^2} \cdot \frac{\partial}{\partial T} (e^{\epsilon/kT})$$

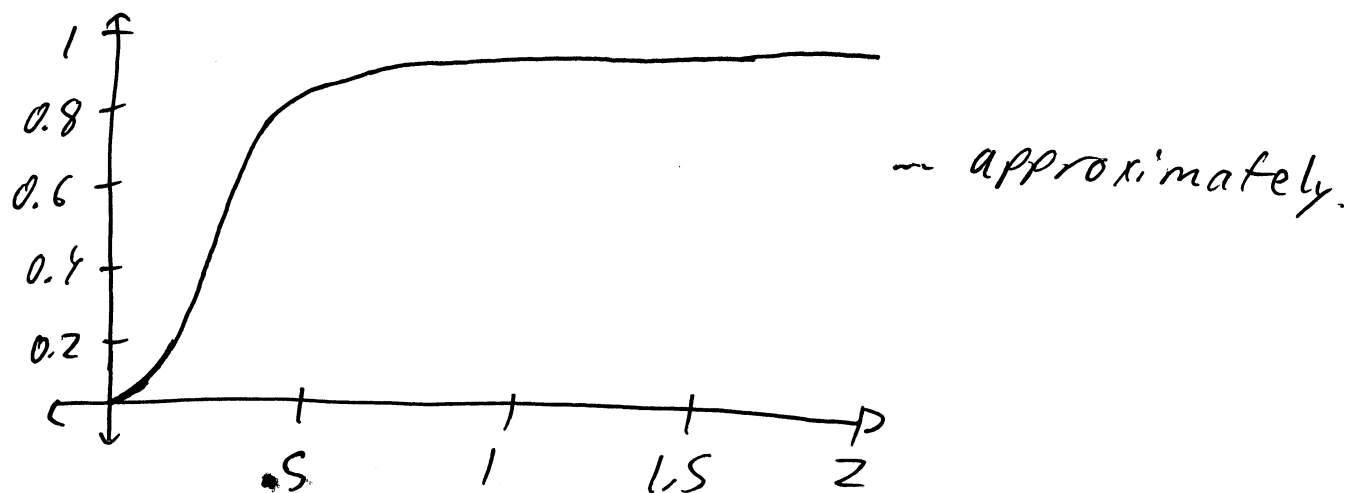
$$C = \frac{N\epsilon^2}{kT^2} \cdot \frac{e^{\epsilon/kT}}{(e^{\epsilon/kT} - 1)^2}$$

d. When $kT \gg \epsilon$, $e^{\epsilon/kT} \approx 1 + \frac{\epsilon}{kT}$

$$C \approx \frac{N\epsilon^2}{kT^2} \frac{\left(1 + \frac{\epsilon}{kT}\right)}{\left(\frac{\epsilon}{kT}\right)^2} \approx \frac{N\epsilon^2}{kT^2} \frac{1}{\left(\frac{\epsilon}{kT}\right)^2} = Nk$$

$C \approx Nk$ Agrees with equipartition theorem, since each oscillator counts as 2 degrees of freedom (1 kinetic, 1 potential).

$$e, \quad \frac{C}{Nk} = \frac{e^{1/T}}{T^2(e^{1/T} - 1)^2}$$



Approximately similar to the curves in Figure 1.14

At high temperatures, the heat capacities are higher than theoretical values because we are measuring C_p instead of C_v .

At low T , the curve predicted by the Einstein model is flatter than the data as it approaches the origin.

C_v reaches half of its equilibrium value at $kT \approx \frac{\epsilon}{3}$. For lead, $T_{1/2} = 22K \Rightarrow$

$\epsilon = .0057 \text{ eV}$. For aluminum, $T_{1/2} = 100K \Rightarrow$

$\epsilon = .026 \text{ eV}$. For diamond, $T_{1/2} = 460 \Rightarrow \epsilon = 0.12 \text{ eV}$

$$7.4 \text{ a. } \Delta S_{\text{body}} = \int_{200}^{400} \frac{1 \text{ kg} \cdot 10 \text{ J/kg} \cdot \text{K}}{T} dT$$

$$= 10 \ln \left(\frac{400}{200} \right) = \boxed{6.93 \text{ J/K}}$$

$$\Delta S_{\text{reservoir}} = \frac{-1 \text{ kg} \cdot 10 \text{ J/kg} \cdot \text{K} \cdot 200 \text{ K}}{400 \text{ K}}$$

$$= \boxed{-5.0 \text{ J/K}}$$

$$b. \Delta S_{\text{body}} = \int_{200}^{100} \frac{10}{T} dT = 10 \ln \left(\frac{100}{200} \right) = \boxed{-6.93 \text{ J/K}}$$

$$\Delta S_{\text{reservoir}} = \frac{1 \cdot 10 \cdot 100 \text{ K}}{100 \text{ K}} = \boxed{10 \text{ J/K}}$$

$$c. \text{ a. } \Delta S_{\text{universe}} = 6.93 - 5.0 = \boxed{1.93 \text{ J/K}}$$

$$b. \Delta S_{\text{universe}} = -6.93 + 10 = \boxed{3.07 \text{ J/K}}$$

7.-8 a. ~~Then~~ $\Delta Q_{net} = 0$

$$m c_p (T_f - T_1) + m c_p (T_f - T_2) = 0$$

$$2 T_f = T_1 + T_2$$

$$T_f = \frac{T_1 + T_2}{2}$$

$$\Delta S_1 = \int_{T_1}^{T_f} \frac{m c_p}{T} dT = m c_p \ln \left(\frac{T_f}{T_1} \right)$$

$$\Delta S_2 = \int_{T_2}^{T_f} \frac{m c_p}{T} dT = m c_p \ln \left(\frac{T_f}{T_2} \right)$$

$$\Delta S_{universe} = m c_p \ln \left(\frac{T_f}{T_1} \right) + m c_p \ln \left(\frac{T_f}{T_2} \right)$$

$$= m c_p \ln \left(\frac{T_f^2}{T_1 T_2} \right) = m c_p \ln \left(\frac{T_f^2}{\sqrt{T_1 T_2}^2} \right)$$

$$= 2 m c_p \ln \left(\frac{T_f}{\sqrt{T_1 T_2}} \right) = \boxed{2 m c_p \ln \left(\frac{T_1 + T_2}{2 \sqrt{T_1 T_2}} \right)}$$

b. True if $T_1 + T_2 > 2 \sqrt{T_1 T_2} \Rightarrow T_1 + T_2 + 2 \sqrt{T_1 T_2}$
 ~~$T_1 + T_2 - 2 \sqrt{T_1 T_2} > 2 \sqrt{T_1 T_2}$~~

$$T_1 - 2\sqrt{T_1 T_2} + T_2 + 2\sqrt{T_1 T_2} > 2\sqrt{T_1 T_2}$$

$$(\sqrt{T_1} - \sqrt{T_2})^2 + 2\sqrt{T_1 T_2} > 2\sqrt{T_1 T_2}$$

Since $(\sqrt{T_1} - \sqrt{T_2})^2 > 0$, this is clearly a true statement.

7-11

$$T ds = c_v dT + T \left(\frac{\partial P}{\partial T} \right)_v dv$$

$$ds = \frac{c_v}{T} dT + \left(\frac{\partial P}{\partial T} \right)_v dv$$

$$\int ds = \int \frac{c_v}{T} dT + \int \left(\frac{\partial P}{\partial T} \right)_v dv$$

$$P = \frac{RT}{v-b} - \frac{a}{v^2}; \quad \left(\frac{\partial P}{\partial T} \right)_v = \frac{R}{v-b}$$

$$S - S_0 = c_v \ln \left(\frac{T}{T_0} \right) + R \ln \left(\frac{v-b}{v_0-b} \right)$$

$$S = c_v \ln T + R \ln(v-b) - \ln(v_0-b) + \underbrace{S_0 - \ln T_0}_{\text{might as well rename these three constants "S}_0\text{"}}$$

$$S = c_v \ln T + R \ln(v-b) + S_0$$

$$7-13 \text{ a. } \kappa = -\frac{1}{V} \left(\frac{\partial V}{\partial P} \right)_T \quad \kappa_s = -\frac{1}{V} \left(\frac{\partial V}{\partial P} \right)_S$$

Setting $ds=0$ in equation (7.12) yields:

$$\left(\frac{\partial V}{\partial P} \right)_S = -\frac{\beta V}{C_p} \cdot \frac{C_v \kappa}{\beta} = -V \kappa \frac{C_v}{C_p}$$

$$\kappa_s = -\frac{1}{V} \cdot -V \kappa \frac{C_v}{C_p} = \kappa \frac{C_v}{C_p}$$

But, from (7.19), we have

$$C_v = C_p - \frac{T_v \beta^2}{\kappa} \implies \frac{C_v}{C_p} = 1 - \frac{T_v \beta^2}{C_p \kappa}$$

$$\kappa_s = \kappa \cdot \frac{C_v}{C_p} = \kappa - \frac{T_v \beta^2}{C_p}$$

Therefore

$$\kappa - \kappa_s = \kappa - \left(\kappa - \frac{T_v \beta^2}{C_p} \right)$$

$$= \frac{T_v \beta^2}{C_p} \text{ as desired.}$$

$$b, \quad p_v = RT \quad p = \frac{RT}{V}$$

For an ideal gas

$$\beta = \frac{1}{V} \left(\frac{\partial V}{\partial T} \right)_p = \frac{1}{V} \left(\frac{R}{p} \right) = \frac{1}{T} ; c_p = \frac{5}{2} R$$

$$\gamma_c - \gamma_s = \frac{T V \cdot \frac{1}{T^2}}{\frac{5}{2} R} = \frac{2}{5} \cdot \frac{V}{RT} = \left(\frac{2}{5} \cdot \frac{1}{p} \right)$$

7-15 a. (Hint Eastwood:

$$\eta = \left(\frac{\partial T}{\partial V} \right)_u = \left(\frac{\partial T}{\partial V} \right)_S + \left(\frac{\partial T}{\partial S} \right)_V \left(\frac{\partial S}{\partial V} \right)_u$$

Setting $ds = 0$ in (7.10) yields

$$\left(\frac{\partial T}{\partial V} \right)_S = - \frac{T\beta}{C_V \kappa}$$

Setting $dV = 0$ in (7.10) yields

$$\left(\frac{\partial T}{\partial S} \right)_V = \frac{T}{C_V}$$

Setting $du = 0$ in (7.9) yields

$$\left(\frac{\partial S}{\partial V} \right)_u = \frac{P}{T}$$

$$\eta = - \frac{T\beta}{C_V \kappa} + \frac{T}{C_V} \cdot \frac{P}{T} = \frac{1}{C_V} \left(P - \frac{T\beta}{\kappa} \right)$$

$$b. \quad u = \left(\frac{\partial T}{\partial P} \right)_h = \left(\frac{\partial T}{\partial P} \right)_S + \left(\frac{\partial T}{\partial S} \right)_P \left(\frac{\partial S}{\partial P} \right)_h$$

Setting $ds = 0$ in (7.11) yields

$$\left(\frac{\partial T}{\partial P} \right)_S = \frac{T_V \beta}{C_P}$$

Setting $dp=0$ in (7.11) yields

$$\left(\frac{\partial T}{\partial s}\right)_p = \frac{T}{c_p}$$

Setting $dh=0$ in the identity

$$Tds = dh - vdp \quad \text{yields}$$

$$\left(\frac{\partial s}{\partial p}\right)_h = -\frac{v}{T}$$

~~Setting $dh=0$ in the identity~~

$$\mu = \frac{Tv\beta}{c_p} - \frac{v}{T} \cdot \frac{T}{c_p}$$

$$\mu = \frac{v}{c_p} (T\beta - 1)$$

C. VdW Gas:

$$\beta = \frac{1}{v} \left(\frac{\partial v}{\partial T}\right)_p \quad \gamma_c = -\frac{1}{v} \left(\frac{\partial v}{\partial p}\right)_T$$

$$p + \frac{a}{v^2} = \frac{RT}{v-b}$$

$$p = \frac{RT}{v-b} - \frac{a}{v^2}$$

Note that $\frac{\beta}{\kappa} = \left(\frac{\partial p}{\partial T} \right)_v$

$$\left(\frac{\partial p}{\partial T} \right)_v = \frac{R}{v-b}$$

So $\eta = \frac{1}{C_v} \left(p - \frac{TR}{v-b} \right)$

To find μ , we still need to compute

$$\left(\frac{\partial v}{\partial T} \right)_p : \left(\frac{\partial v}{\partial T} \right)_p = - \frac{\left(\frac{\partial p}{\partial T} \right)_v}{\left(\frac{\partial p}{\partial v} \right)_T} \quad (\text{Cyclic relation})$$

$$= \frac{-\frac{R}{v-b}}{-\frac{RT}{(v-b)^2} + \frac{2a}{v^3}} = \frac{R(v-b)}{RT - \frac{2a(v-b)^2}{v^3}}$$

$$\left(\frac{\partial v}{\partial T} \right)_p = \frac{R(v-b)v^3}{RTv^3 - 2a(v-b)^2}$$

$$\beta = \frac{R(v-b)v^2}{RTv^3 - 2a(v-b)^2}$$

$$u = \frac{v}{c_p} \left(\frac{TR(v-b)v^2}{RTv^3 - 2a(v-b)^2} - 1 \right)$$

For an ideal gas, $\beta = \frac{1}{T}$ & $\kappa = \frac{1}{p}$

$$\eta = \frac{1}{c_v} \left(p - T \cdot \frac{p}{T} \right) = 0 \checkmark$$

$$u = \frac{v}{c_p} \left(T \cdot \frac{1}{T} - 1 \right) = 0 \checkmark$$