Show work to get credit. There are problems where the answer is a specific number.

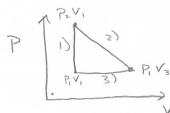
(1) (5 pts) 2 grams of ice at 0° C is melted to 2 grams of water at 0° C. This occurs at 1 atm of pressure. a) What is the change of entropy of this system? b) Did the number of states of this system increase or decrease? c) Determine the ratio (final number of states)/(initial

a) 
$$dS = \frac{666J}{273K} = 2.44 = \frac{1}{K}$$

b) From Eq 2. 45 Increase because 
$$\mathcal{N} = e^{\frac{5}{128}}$$
 and Sincrease c)  $\frac{\mathcal{R}_{fin}}{\mathcal{R}_{init}} = e^{\frac{dS}{l_{BB}}} = e^{\frac{2.44 \frac{7}{l_{K}}}{1.38 \times 10^{-23} \frac{7}{l_{K}}}} = e^{\frac{1.77 \times 10^{23}}{1.38 \times 10^{-23} \frac{7}{l_{K}}}} = e^{\frac{1.77 \times 10^{23}}{1.38 \times 10^{-23} \frac{7}{l_{K}}}}$ 

c) 
$$\frac{\mathcal{R}_{fin}}{\mathcal{R}_{init}} = e^{dS/l_{eB}} = e^{\frac{2.44 \frac{T_{k}}{T_{k}}}{1.38 \times 10^{-23 \frac{T_{k}}{T_{k}}}}} = e^{1.77 \times 10^{23}}$$

(2) (5 pts) You make a heat engine which has He gas in a piston. It goes through 3 steps. 1) Starting at  $P_1$ ,  $V_1$  the pressure is increased to  $P_2$  keeping the volume fixed. 2) Then the volume is expanded until the pressure is back to  $P_1$  but the volume is now  $V_3$ . For this step the pressure is linearly dependent on V. 3) Then the volume is decreased back to  $V_1$  keeping the pressure constant. a) Draw this cycle on a PV diagram; label each step and the various  $P_i, V_i$  points. b) Make a table that has the amount of heat added to the gas and the work done on the gas for each step. c) Compute the total work performed in one cycle and the The has 3 guard, D.O.F. total heat added to the gas in one cycle.



$$\frac{1}{2} \left( \frac{1}{2} \left( \frac{1}{2} - \frac{1}{2} \right) \right) \left( \frac{3}{2} \left( \frac{1}{2} - \frac{1}{2} \right) \right) \left( \frac{3}{2} \left( \frac{1}{2} - \frac{1}{2} \right) \right) + \left( \frac{1}{2} + \frac{1}{2} \right) \left( \frac{1}{2} - \frac{1}{2} \right) \left( \frac{3}{2} \left( \frac{1}{2} - \frac{1}{2} \right) \right) + \left( \frac{1}{2} + \frac{1}{2} \right) \left( \frac{1}{2} - \frac{1}{2} \right) \left( \frac{3}{2} - \frac{1}{2} \right) \left( \frac{3}{$$

$$W = -\int_{v_i}^{v_i} P dv \qquad \Delta U = \frac{3}{2} N k_B \Delta T = \frac{3}{2} (R_F V_i - R_F V_i)$$

$$Q = \Delta U - W$$

$$V \text{ on gas}$$

$$W_{TOT} = (V_3 - V_1)[P_1 - \frac{P_2 + P_1}{2}] = -\frac{1}{2}(V_3 - V_1)(P_2 - P_1)$$

$$W_{performed} = -W_{TOT} = \frac{1}{2}(V_3 - V_1)(P_2 - P_1)$$

$$Q = \frac{3}{2}(P_2 - P_1)V_1 + \frac{3}{2}(P_1V_3 - P_2V_1) + \frac{3}{2}(P_1V_1 - P_1V_3)$$

$$-W_{TOT}$$

$$Q = -W_{TOT} = \frac{1}{2}(V_3 - V_1)(P_2 - P_1)$$

a) ?rab 5.2 b) ?rab 5.2

(3) (5 pts) Consider the reaction of C (graphite) plus  $2H_2$  going to  $CH_4$  that takes place at standard temperature and pressure. a) Specify how much energy *you* have to add or remove to make this reaction go for each mole of  $CH_4$ . b) For this reaction, verify that the data for the Gibbs free energy, enthalpy, and entropy are consistent.

Need to check  $\Delta G$  for the left and right side  $C+2H_2 \rightarrow CH_y$   $E_{you} = \Delta G_{CH_y} - \Delta G_c - 2\Delta G_{H_2} = -50,72 \text{ kJ} - 0 - 0 = -50.72 \text{ kJ}$ You set out 50.72 kJ

b) From Eg (5.9)

ΔG = ΔH - TΔ5

-50.72 ×10<sup>3</sup> J = -74.81 ×10<sup>3</sup> J - 298 κ (.186.26 = -5.74 = -2×130.68)

= -74.81 kJ + 24.09 kJ

-50.72 kJ matches

(4) (5 pts) For this problem, I want all answers to 5 significant digits. Snedley Frumble has discovered a quantum system where the energy levels are  $E_n = \varepsilon n^3$  with n = 0, 1, 2, ... and there is one state with n = 0, two states with n = 1, three states with n = 2, etc. For the case  $\varepsilon/(k_BT) = \ln(5/4)$ , a) compute the probability to have energy  $\varepsilon$ .

$$Z = 1 \cdot 1 + 2 \cdot e^{-\beta \xi} + 3 \left(e^{-\beta \xi}\right)^{8} + \dots$$

$$= 1 + 2 + 3 \left(\frac{4}{5}\right)^{8} + 4 \left(\frac{4}{5}\right)^{27} + 5 \left(\frac{4}{5}\right)^{64} = 3.11299$$

b) 
$$P_{E=E} = \frac{2.4/5}{Z} = 0.51398$$

(5) (10 pts) Aluminum silicate, Al<sub>2</sub>SiO<sub>5</sub>, has three different crystalline forms: kyanite, andalusite, and sillimanite. a) Which is the stable form at T = 298 K and 1 bar? Explain why you chose that form. b) As you raise the pressure keeping T = 298 K, are there phase changes? If there are phase changes, determine all of the pressures where they occur. c) Instead of raising the pressure, you raise the temperature keeping P=1 bar. Are there phase changes? If there are phase changes, determine all of the temperatures where they

a) Find the smallest DG Kyanite

b) Use dG = VdP

The phase changes occur when

DGa + VAP = DG' + V, AP

 $\Delta P = \frac{\Delta G_a - \Delta G_b}{V_b - V_a}$ 

From the table Vandal. > Vkyan and Vsilli, > Vkyan

This means DG kyan + Vkyan DP is always less than that

for the other forms => No Phase Transition

c) Use dG = - SdT

The phase changes occur when DG - SaDT = DG - SDT

ΔT = ΔG° - ΔG° Sa - SL

andal. DT = 1.722×1035 = 130K => T= 428K = lower means

AT = 2.89×1035 = 235k => T= 533k | kyamite -> andalusite

Now check whether andalusite -> sillimanite

 $\Delta T = \frac{1.67 \times 10^3 \text{J}}{2.89 \text{J}} = 578 \text{K} = 3 T = 876 \text{K}$ 

andalusite -> sillimanite at 876K

Sec. 3.3

(6) (10 pts) A system consists of  $N \gg 1$  quantum objects each of which has two two states with energies:  $E_1 = 0$  and  $E_2 = \varepsilon$ . Define  $N_1$  to be the number of objects in state 1 and  $N_2$  to be the number of objects in state 2. This system is isolated. It is known to have internal energy  $U = N_2 \varepsilon$ . a) Determine the entropy for this system in terms of U and N. b) Determine the chemical potential in terms of U and N.

temperature and

Use 
$$S = k_B \ln \mathcal{R}$$
  
For  $Z$  state  $Sy$  stem  $\mathcal{R} = \frac{N!}{N_2! (N-N_2)!}$   
 $S = k_B \ln \frac{N!}{N_2! (N-N_2)!}$   
 $= k_B \ln \frac{(\frac{N}{\epsilon})^N}{(\frac{N-N_2}{\epsilon})^{N-N_2}} \int \frac{St_1 r ling}{St_1 r ling} \frac{approx}{approx}$   
 $= k_B [N \ln N - N_2 \ln N_2 - (N-N_2) \ln (N-N_2)]$   $= k_B [N \ln N - \frac{U}{\epsilon} \ln (\frac{U}{\epsilon}) - (N-\frac{U}{\epsilon}) \ln (N-\frac{U}{\epsilon})]$   
 $= \frac{1}{2} \ln \frac{S}{s} \ln (\frac{N-U}{s}) \ln (N-\frac{U}{s}) + \frac{k_B}{\epsilon} = \frac{1}{2} \ln (\frac{N-U}{s})$   
 $= \frac{k_B}{s} \ln (\frac{N-U}{s})$ 

$$M = -T \left( \frac{\partial S}{\partial N} \right)_{U,V} = -k_B T \left( lm N + 1 - lm \left( N - \frac{U}{E} \right) - 1 \right) \qquad E_{g.(3.59)}$$

$$= k_B T lm \left( \frac{N E - U}{N E} \right)$$

$$= E lm \left( \frac{N E - U}{N E} \right)$$

$$= lm \left( \frac{N E - U}{N} \right)$$