## PHYSICS 140A MIDTERM EXAM 2

Tuesday November 10, 2009

$$T \ dS = C_V \ dT + T \frac{\beta}{\kappa} \ dV$$
$$T \ dS = C_p \ dT - VT \ \beta \ dp$$
$$T \ dS = C_V \frac{\kappa}{\beta} \ dp + \frac{C_p}{\beta V} \ dV$$

1. In this problem, we consider 3 heat engines. The 3 cycles of the working substance are drawn in the S-T plane below. Cycle 1 is ABEHA; cycle 2 is ABCDEHA; and cycle 3 is ABEFGHA. Assume all 3 cycles are reversible. Also, assume T'' < T' < T and S'' < S' < S.

(a) What is the efficiency  $\eta_1$  of heat engine 1?

(b) What is the efficiency  $\eta_2$  of heat engine 2?

- (c) What is the efficiency  $\eta_3$  of heat engine 3?
- (d) Compare the 3 efficiencies  $\eta_1$ ,  $\eta_2$  and  $\eta_3$ .

(e) Since the cycles are reversible, we can consider the combination consisting of cycle 3 and the reverse of cycle 1. What is the efficiency  $\eta$  of this new cycle? What can one say about the efficiencies  $\eta_1$ ,  $\eta_3$  and  $\eta$  relative to one another?

(a)

$$\eta_1 = 1 - \frac{T}{T}$$

(b) 
$$\eta_2 = 1 - \frac{T'}{T}$$

(c) 
$$\eta_3 = 1 - \frac{T^{\prime\prime}}{T}$$

(d) 
$$n_3 > n_1 =$$

(e) n = 1

$$\eta = 1 - \frac{T''}{T'}$$
$$\eta_3 > \eta_1$$
$$\eta_3 > \eta$$

 $\eta_2$ 

One cannot determine whether  $\eta_1$  or  $\eta$  is larger without more information about relative sizes of ratios T'/Tand T''/T' 2. Consider the reversible, closed cycle of an ideal gas, which consists of

- (i) an adiabatic expansion from state A to state B
- (ii) an isobaric compression from state B to state C
- (iii) an isochoric transformation from state C to state A

The cycle is drawn in the V-p plane below. The coordinates of states A, B and C are  $(V_A, p_A)$ ,  $(V_B, p_B)$ , and  $(V_A, p_B)$ , respectively. Assume that the heat capacity of the ideal gas at constant volume is  $C_V = \alpha nR$ , for  $\alpha$  some constant.

(a) Compute the change in entropy S for each transformation of the cycle. (Compute  $S_{BA}$ ,  $S_{CB}$  and  $S_{AC}$ .) Check your result by verifying that the total change in entropy around the closed cycle is what you expect.

(b) Compute the heat Q for each transformation of the cycle. (Compute  $Q_{BA}$ ,  $Q_{CB}$  and  $Q_{AC}$ .) What is the total heat  $Q_{TOT}$  for the cycle?

(c) Compute the work W for each transformation of the cycle. (Compute  $W_{BA}$ ,  $W_{CB}$  and  $W_{AC}$ .) What is the total work  $W_{TOT}$  for the cycle?

(d) Compare  $Q_{\text{TOT}}$  and  $W_{\text{TOT}}$ ? What do you conclude about the change in internal energy U for the cycle?

(a)

$$T \ dS = C_V \frac{\kappa}{\beta} \ dp + \frac{C_p}{\beta V} \ dV$$

For ideal gas,  $C_V = \alpha n R$ ,  $C_p = (\alpha + 1)nR$ ,  $\beta = \frac{1}{T}$ ,  $\kappa = \frac{1}{p}$ , pV = nRT.

$$T \ dS = \alpha V \ dp + (\alpha + 1)p \ dV$$

$$dS = \alpha nR \frac{dp}{p} + (\alpha + 1)nR \frac{dV}{V}$$
$$S_{BA} = 0$$
$$S_{CB} = (\alpha + 1) nR \ln\left(\frac{V_A}{V_B}\right)$$
$$S_{AC} = \alpha nR \ln\left(\frac{p_A}{p_B}\right)$$

$$S_{\text{TOT}} = (\alpha + 1) \ nR \ \ln\left(\frac{V_A}{V_B}\right) + \alpha \ nR \ \ln\left(\frac{p_A}{p_B}\right)$$

 $A \to B$  is an adiabatic expansion, so

$$p_A V_A^{\gamma} = p_B V_B^{\gamma}$$
$$\frac{p_A}{p_B} = \left(\frac{V_B}{V_A}\right)^{\frac{\alpha+1}{\alpha}}$$

Thus,

$$S_{\text{TOT}} = (\alpha + 1) \ nR \ \ln\left(\frac{V_A}{V_B}\right) + \alpha \ nR \ \ln\left(\frac{V_B}{V_A}\right)^{\frac{\alpha+1}{\alpha}} = 0$$

as expected.

(b)

$$dQ_{\rm rev} = T \ dS = \alpha V \ dp + (\alpha + 1)p \ dV$$
 
$$Q_{BA} = 0$$

for adiabatic transformation.

$$Q_{CB} = (\alpha + 1) \ p_B \int_{V_B}^{V_A} dV = (\alpha + 1) \ p_B (V_A - V_B) < 0$$
$$Q_{AC} = \alpha V_A \int_{p_B}^{p_A} dp = \alpha V_A (p_A - p_B) > 0$$
$$Q_{\text{TOT}} = (\alpha + 1) \ p_B (V_A - V_B) + \alpha V_A (p_A - p_B)$$

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$$dW_{\rm rev} = pdV$$

$$W_{BA} = \frac{1}{(1-\gamma)} \left( p_B V_B - p_A V_A \right) = \alpha \left( p_A V_A - p_B V_B \right) > 0$$
since  $\gamma = \frac{\alpha+1}{\alpha} \Rightarrow \frac{1}{1-\gamma} = -\alpha$ 

$$W_{CB} = p_B \left( V_A - V_B \right)$$

$$W_{AC} = 0$$

$$W_{\rm TOT} = \alpha p_A V_A - (\alpha+1) p_B V_B + p_B V_A$$

(d)

$$Q_{\rm TOT} = W_{\rm TOT}$$

so  $U_{\rm TOT}=0$  for the closed cycle as expected.

3. Short problems.

(a) Prove for a general thermodynamic system that

$$\left(\frac{\partial S}{\partial V}\right)_T = \left(\frac{\partial p}{\partial T}\right)_V.$$

(b) If two thermodynamic systems A and B are in thermodynamic equilibrium, what do you know about them?

(c) A liquid consists of a mixture of 3 different liquids in thermodynamic equilibrium. How many independent variables are required to describe the system? List the independent variables.

(d) The solid and liquid phases of a given substance are in thermodynamic equilibrium. What do you know about the specific Gibbs free energies of the solid and the liquid?

(a)

$$\begin{split} S &= -\left(\frac{\partial F}{\partial T}\right)_V \\ p &= -\left(\frac{\partial F}{\partial V}\right)_T \\ \left(\frac{\partial S}{\partial V}\right)_T &= -\frac{\partial^2 F}{\partial V \partial T} = -\frac{\partial^2 F}{\partial T \partial V} = \left(\frac{\partial p}{\partial T}\right)_V \end{split}$$

(b)

(d)

$$T_A = T_B$$
$$p_A = p_B$$
$$\mu_A = \mu_B$$

(c) k = 3 and  $\pi = 1$  $f = k - \pi + 2 = 3 - 1 + 2 = 4$ 

Independent variables: T, p, and 2 different mole fractions  $x_1 = \frac{n_1}{n_1 + n_2 + n_3}$ ,  $x_2 = \frac{n_2}{n_1 + n_2 + n_3}$ Note that  $x_3 = 1 - x_1 - x_2$ , so one of the three mole fractions is not independent.

The specific Gibbs free energies of the solid and liquid phases are equal.

4. A gas (NOT an ideal gas) has Helmholtz free energy

$$F(T, V, n) = -nRT \left\{ \ln T^{3/2} + \ln V - \ln n + 1 + c \right\} + n^2 RT \frac{B(T)}{V}$$

where c is a constant and B(T) is an arbitrary function of temperature T.

- (a) Find the entropy S of the gas.
- (b) Find the pressure p of the gas.
- (c) Find the chemical potential  $\mu$  of the gas.
- (d) Find  $C_V$ , the heat capacity at constant volume.
- (e) Find the internal energy U of the gas.
- (f) Find the Gibbs free energy G of the gas.

 $S = -\left(\frac{\partial F}{\partial T}\right)_{V,n}$  $S = nR\left\{\ln T^{3/2} + \ln V - \ln n + 1 + c\right\} + \frac{3}{2}nR - n^2R\frac{B(T)}{V} - n^2RT\frac{B'(T)}{V}$ 

(b)

(a)

$$p = -\left(\frac{\partial F}{\partial V}\right)_{T,n}$$
$$p = \frac{nRT}{V} + \frac{n^2 RTB(T)}{V^2}$$
$$p = \frac{nRT}{V} \left(1 + \frac{nB(T)}{V}\right)$$

(c)

$$\mu = \left(\frac{\partial F}{\partial n}\right)_{T,V}$$
$$\mu = -RT\left\{\ln T^{3/2} + \ln V - \ln n + 1 + c\right\} + RT + 2nRT\frac{B(T)}{V}$$
$$\mu = -RT\left\{\ln T^{3/2} + \ln V - \ln n + c\right\} + 2nRT\frac{B(T)}{V}$$

(d)

$$C_{V} = T \left(\frac{\partial S}{\partial T}\right)_{V,n}$$
$$\left(\frac{\partial S}{\partial T}\right)_{V,n} = \frac{3nR}{2T} - 2n^{2}R\frac{B'(T)}{V} - n^{2}RT\frac{B''(T)}{V}$$
$$C_{V} = T \left(\frac{\partial S}{\partial T}\right)_{V,n} = \frac{3}{2}nR - 2n^{2}RT\frac{B'(T)}{V} - n^{2}RT^{2}\frac{B''(T)}{V}$$
$$C_{V} = \left(\frac{\partial U}{\partial T}\right)_{V,n}$$

$$U = F + TS$$
$$U = \frac{3}{2}nRT - n^2RT^2\frac{B'(T)}{V}$$

(f)

$$\begin{split} G &= F + pV \\ G &= -nRT \left\{ \ln T^{3/2} + \ln V - \ln n + 1 + c \right\} + n^2 RT \frac{B(T)}{V} + nRT + \frac{n^2 RTB(T)}{V} \\ G &= -nRT \left\{ \ln T^{3/2} + \ln V - \ln n + c \right\} + 2n^2 RT \frac{B(T)}{V} \\ G &= \mu n \end{split}$$