

<b>Q1</b>	<b>/15</b>
<b>Q2</b>	<b>/15</b>
<b>Q3</b>	<b>/16</b>
<b>Q4</b>	<b>/15</b>
<b>Q5</b>	<b>/15</b>
<b>Q6</b>	<b>/20</b>
<b>Q7</b>	<b>/15</b>
<b>Q8</b>	<b>/10</b>
	<b>/121</b>

UNIVERSITY OF TORONTO  
FACULTY OF APPLIED SCIENCE AND ENGINEERING

**FINAL EXAMINATION**

APRIL 2013

First Year

APS 104S

INTRODUCTION TO MATERIALS AND CHEMISTRY

Exam Type B

Examiners: *T. Bender, N. P. Kherani, T. Mirkovic*

Duration: 2 hour 30 minutes

**NAME:** \_\_\_\_\_  
 Last \_\_\_\_\_ First \_\_\_\_\_

**STUDENT NO:** \_\_\_\_\_

**INSTRUCTIONS:**

- This is a Type B examination. Only non-programmable calculators are allowed.
- Answer all 8 questions.
- All work is to be done on the pages of this booklet.
- When answering the questions **include all the steps** of your work on these pages and then **write the answer in** the respective **boxes** where provided. For additional space, you may use the back of the preceding page.
- Do not unstaple this exam booklet.
- A Formula Sheet and the periodic table are attached to the end of this exam booklet; if you wish, you may tear-off these sheets *only*.

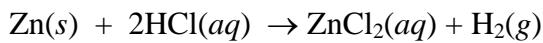
Last Name:

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Student No.

**QUESTION 1** [15 marks]

**Part I.** A sample of zinc metal reacts completely with an excess of hydrochloric acid:



The reaction is run at 273 K and 1.0 atm.

(a) [4 marks] Calculate the work done by the reaction if 19.8 g of zinc is reacted.

(Neglect volume changes other than the change in gas volume)

(b) [5 marks] If the above reaction is exothermic, determine if the following are positive, negative, or zero (circle the correct response) :

$\Delta U$ :	+ve	-ve	0
$\Delta H$ :	+ve	-ve	0
$q$ :	+ve	-ve	0
$\Delta S$ :	+ve	-ve	0
$\Delta G$ :	+ve	-ve	0

(c) [2 marks] In a second experiment, the reaction in (a) is run at 400 K. Which statement is correct?

- i) The system is doing more work at 400 K than at 273 K.
- ii) The system is doing less work at 400 K than at 273 K.
- iii) The surroundings are doing more work at 400 K than at 273 K.
- iv) The surroundings are doing less work at 400 K than at 273 K.

**Part II.**

[4 marks]

(a) A balloon contains 0.76 mol He, 0.18 mol Ne, 0.031 mol Ar and 0.026 mol Rn at 749 mm Hg. What is the partial pressure of Ne?

(b) Which of the noble gases in the above mixture should show the greatest deviation from the ideal gas law at high pressures. Circle the correct answer.

He

Ne

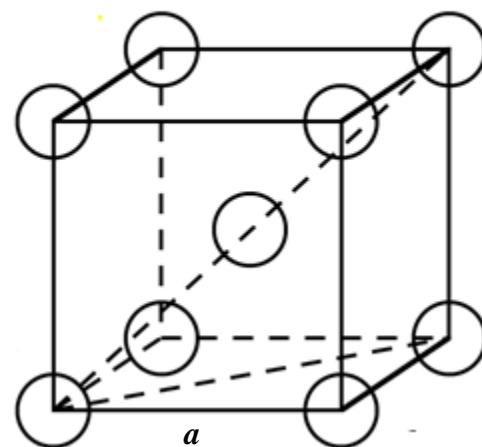
Ar

Rn

all deviate equally

**QUESTION 2 [15 marks]**

- (a) Consider the body centered cubic crystal structure (shown in the small sphere representation on the right) with a unit cell edge length of  $a$  and atomic radius  $R$ . Derive the expression for the lattice constant  $a$  in terms of the atomic radius  $R$ . [3 marks]



- (b) Calculate the atomic packing factor for the BCC crystal structure. [4 marks]

- (c) Calculate the linear atomic density in the close-packed direction of the BCC crystal structure. [3 marks]

- (d) Calculate the atomic radius of iron given that it has a BCC crystal structure, an atomic weight of 55.85 g/mole and mass density of 7.87 g/cm<sup>3</sup>. [5 marks]

Last Name:

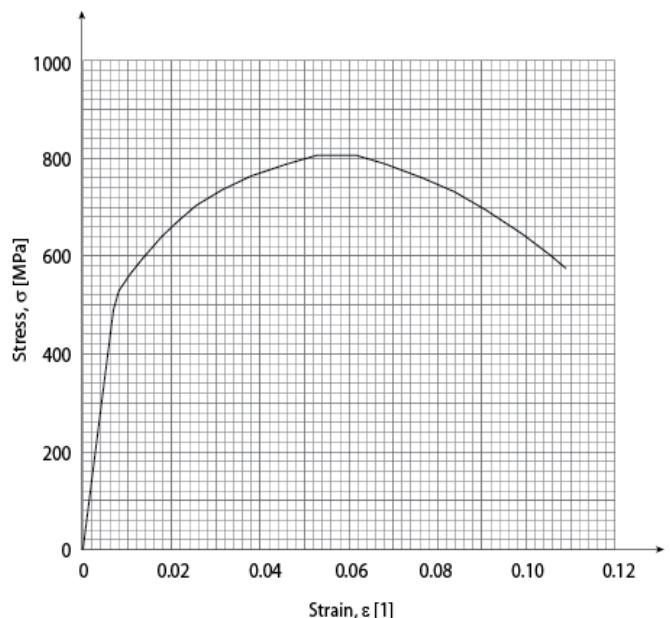
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Student No.

**QUESTION 3** [16 marks]

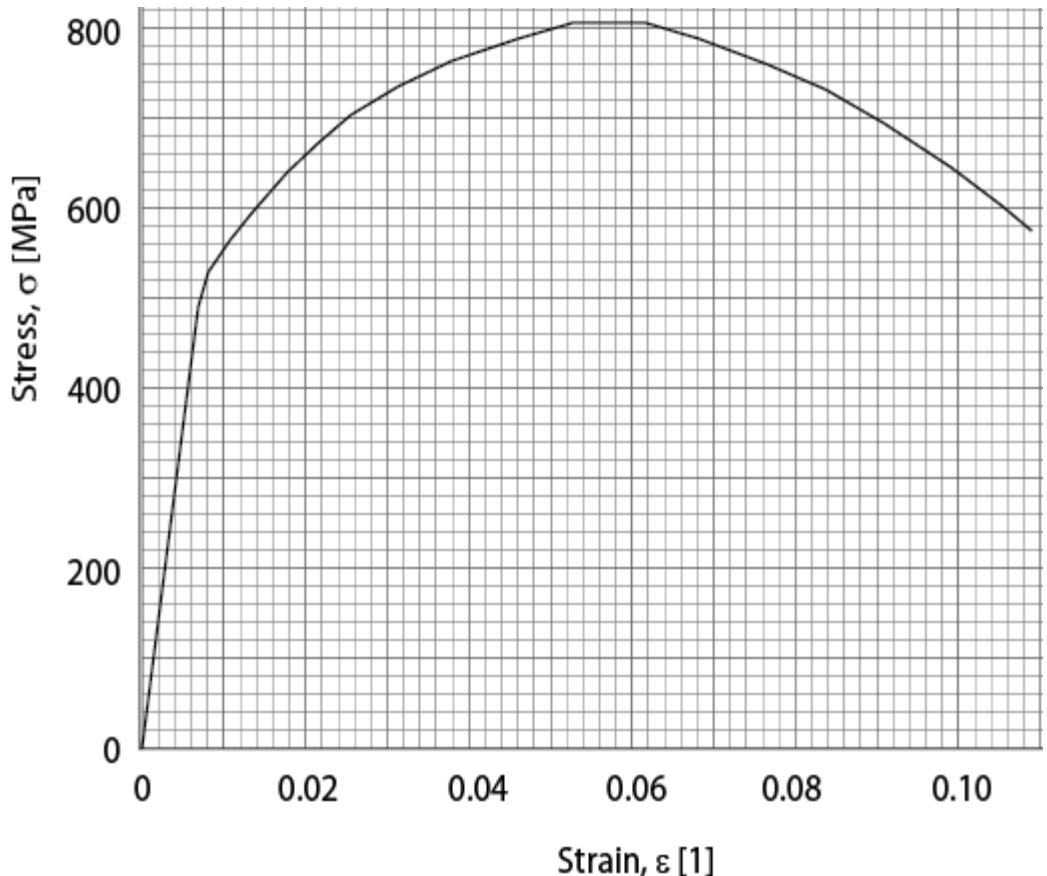
- (i.) Consider the engineering stress-strain curve for a particular alloy shown on the right. Clearly mark the following features/points on the graph on the right: [6 marks]

- A. Tensile Strength.
- B. Fracture Strength.
- C. Proportional Point.
- D. Shade the region that represents resiliency.
- E. Highlight the strain range that represents 'strain hardening'.
- F. Mark the point where 'necking' starts.



- (ii.) Using the expanded engineering stress-strain curve shown below, calculate:

- 1) Young's modulus.  
[2 marks]



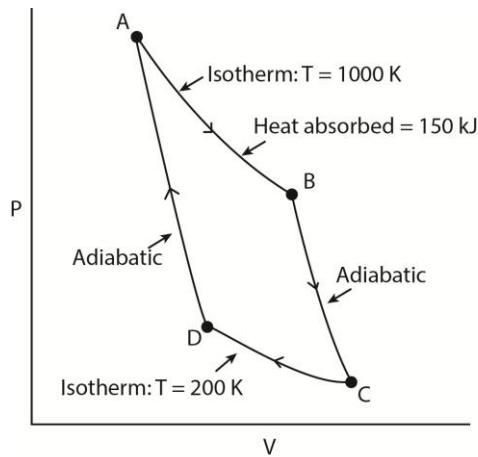
- 2) 0.5% off-set yield strength. [2 marks]

- 3) Strain at fracture.  
[2 marks]

- 4) Given a Poisson coefficient of 0.3, original specimen length of 100 mm, original specimen diameter of 10 mm, and applied tensile stress of 400 MPa, calculate the elongated length and the corresponding diameter of the specimen. [4 marks]

**QUESTION 4** [15 marks]

The diagram below represents a reversible Carnot cycle for an ideal gas:



(a) [2 marks] What is the thermodynamic efficiency of the engine?

(b) [2 marks] How much heat is rejected at the lower temperature, 200 K, during the isothermal compression?

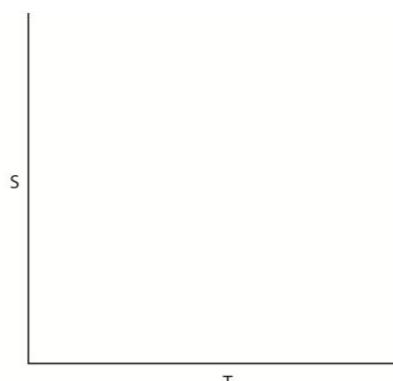
(c) [2 marks] What is the entropy increase during the isothermal expansion at 1000 K?

(d) [2 marks] What is the entropy decrease during the isothermal compression at 200 K?

(e) [2 marks] What is the overall entropy change for the entire cycle?

(f) [2 marks] What is the change in Gibbs free energy during the process A → B?

(g) [3 marks] The above diagram shows a Carnot cycle in the form of a pressure-volume diagram. Sketch the corresponding entropy-temperature diagram, labeling the individual steps A → B (isothermal), B → C (adiabatic), C → D (isothermal), and D → A (adiabatic).



Last Name:

First Name:

Student No.

**QUESTION 5** [15 marks]

If the reaction  $\text{NO}_2(\text{g}) \rightleftharpoons \text{NO}(\text{g}) + \frac{1}{2}\text{O}_2(\text{g})$  comes to equilibrium at a total pressure of 1 bar, analysis of the gas shows that at 700 K,  $\frac{P_{\text{NO}}}{P_{\text{NO}_2}} = 0.872$  and  $\frac{P_{\text{O}_2}}{P_{\text{NO}}} = 0.500$ , and at 800 K,  $\frac{P_{\text{NO}}}{P_{\text{NO}_2}} = 2.50$  and  $\frac{P_{\text{O}_2}}{P_{\text{NO}}} = 0.500$ , if only  $\text{NO}_2(\text{g})$  was initially present in the gas phase.

- (a) [6 marks] Calculate  $K_p$  at 700 and 800 K.

- (b) [3 marks] Calculate  $\Delta H_{\text{reaction}}^\circ$  assuming that it is independent of temperature.

Last Name:

First Name:

Student No.

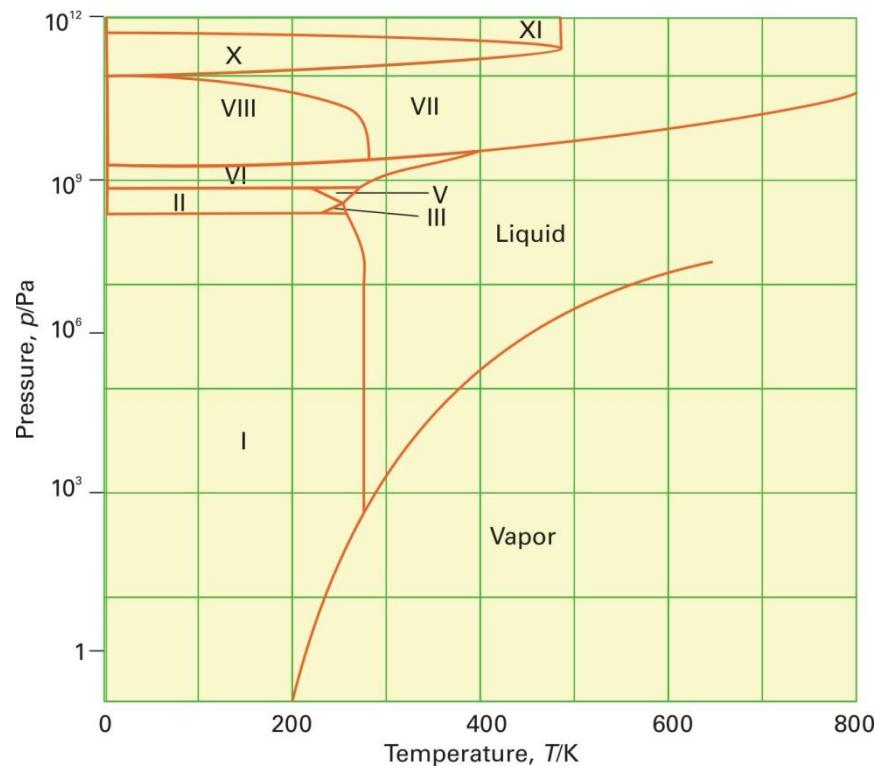
(c) [3 marks] Calculate  $\Delta G_{reaction}^\circ$  for this reaction at 298.15 K.

(d) [3 marks] How would the equilibrium position change, if the following actions were performed on this reaction system? Circle the correct response.

- |   |                      |                     |                      |
|---|----------------------|---------------------|----------------------|
| i) increase in temperature:             | <i>move to right</i> | <i>move to left</i> | <i>doesn't shift</i> |
| ii) decrease the reaction flask volume: | <i>move to right</i> | <i>move to left</i> | <i>doesn't shift</i> |
| iii) addition of inert gas:             | <i>move to right</i> | <i>move to left</i> | <i>doesn't shift</i> |

**QUESTION 6** [20 marks]

6-1. Consider the following phase diagram of water:



(a) Circle on the phase diagram the triple point and explain below what you would observe if the system was at the triple point. [2 marks]

(b) Circle on the phase diagram the critical point. State below the significance of the critical point. [2 marks]

(c) Explain below what happens as you move along a constant pressure line at  $10^6 \text{ Pa}$  starting from 0 K and moving to 600K. [2 marks]

**6-2.** Consider the following phase diagram:

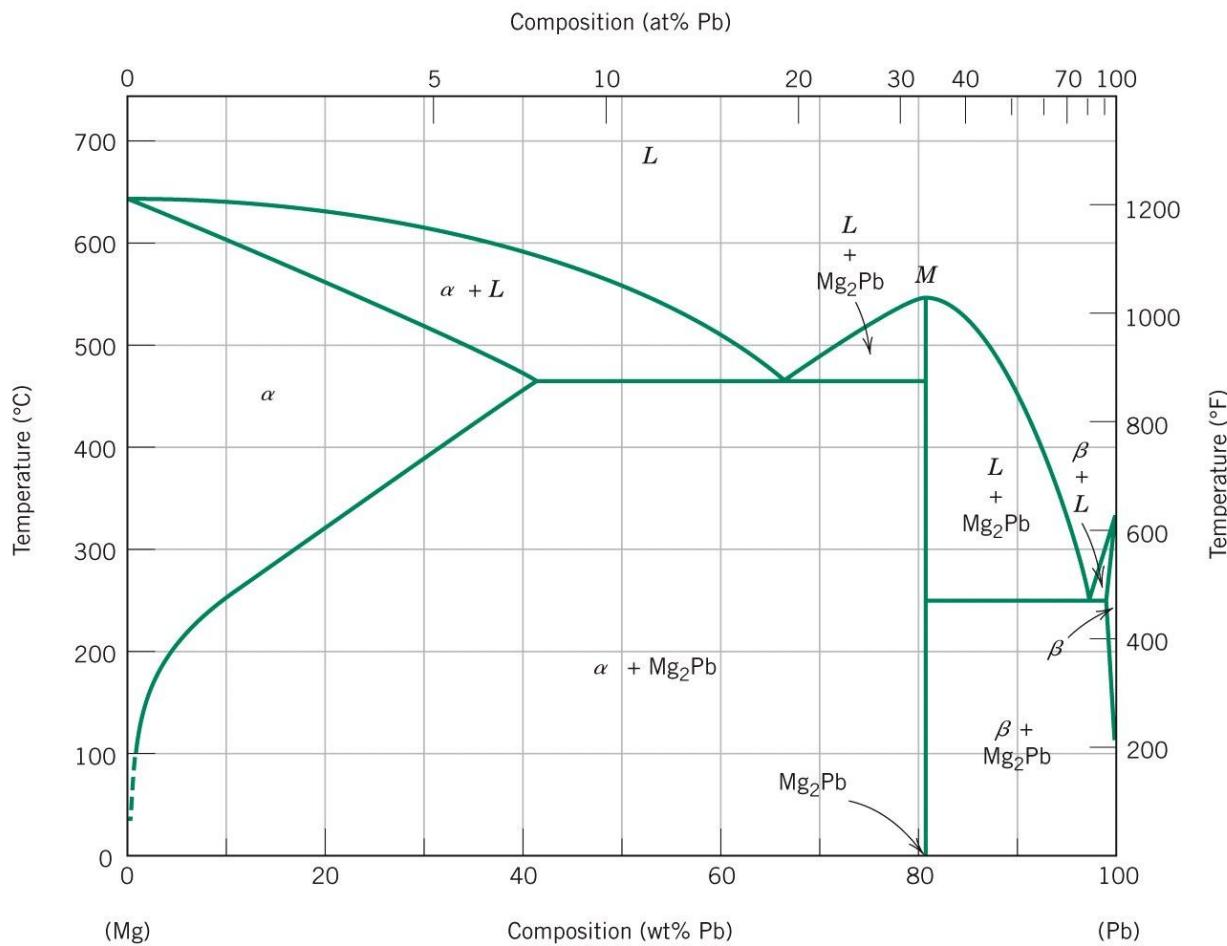


Figure 10.20  
© John Wiley & Sons, Inc. All rights reserved.

(a) For a mixture of 37 kg of Pb and 6.5 kg of Mg state the phase(s) that are present at 800 °C. [2 marks]

(b) Does Pb-Mg form a eutectic mixture? If yes: circle a eutectic point on the phase diagram and identify a eutectic isotherm. If no: explain why with reference to the phase diagram. [2 marks]

(c) Does Pb-Mg form a stoichiometric compound? If yes: identify it and state its melting point? If no: explain why with reference to the phase diagram. [2 marks]

Last Name:

First Name:

Student No.

Parts d-g pertain to the following:

A 50wt% Pb-50wt% Mg mixture is slowly cooled from 700 °C to 300 °C.

(d) At what temperature does the first solid phase form? [1 mark]

Answer:

(e) What is the composition of the first small amount of solid phase formed? [3 marks]

Answer:

(f) At what temperature does the liquid completely solidify? [1 mark]

Answer:

(g) What is the composition of the last remaining liquid phase just before complete solidification? [3 marks]

Answer:

Last Name:

First Name:

Student No.

**QUESTION 7 [15 marks]**

A voltaic electrochemical cell consists of a silver electrode in contact with 346 mL of 0.100 M AgNO<sub>3</sub> solution and a magnesium electrode in contact with 288 mL of a 0.100 M Mg(NO<sub>3</sub>)<sub>2</sub> solution.



(a) [2 marks] Write the overall reaction and the shorthand notation for this voltaic cell.

(b) [3 marks] Calculate the value of  $E$  for the cell at 25°C.

(c) [6 marks] A current is drawn from the cell until 1.20 g of silver have been deposited at the silver electrode. Calculate the value of  $E$  for the cell at this stage of operation.

(d) [1 mark] Determine  $E_{\text{cell}}$  once the cell has reached equilibrium.

(e) [3 marks] True or False? Circle the correct response.

(i) In the electrolyte of an electrochemical cell, current is carried by anions moving toward the anode and cations moving in the opposite direction.      T    or    F

(ii) Electrons are produced at the cathode of a voltaic cell.      T    or    F

(iii) In a fuel cell, an external source of electrical power is used to drive a non-spontaneous reaction in which a fuel is produced.      T    or    F

**QUESTION 8** (10 marks)

(a) Draw the energy band diagrams for an insulator, semiconductor, and metal – making sure to clearly highlight the differences. [3 marks]

Insulator

Semiconductor

Metal

(b) Calculate the number of free electrons per cubic meter of silver, assuming there are 1.5 free electrons per atom of silver. The electrical conductivity, density and atomic weight of silver are  $6.8 \times 10^7 (\Omega \cdot m)^{-1}$ ,  $10.49 \text{ g cm}^{-3}$  and  $107.87 \text{ g mol}^{-1}$  respectively. [3 marks]

Answer:

(c) Using the results from part (b) compute the electron mobility for silver. [3 marks]

Answer:

(d) Compute the resistance of a copper wire 3mm in diameter and 2 m in length given the conductivity of copper is  $6.0 \times 10^7 (\Omega \cdot m)^{-1}$ . [1 mark]

Answer:

Last Name: \_\_\_\_\_ First Name: \_\_\_\_\_ Student No. \_\_\_\_\_

**FORMULAE & CONSTANTS** (You may tear this sheet off.)

$$R = 8.3145 \text{ J}\cdot\text{K}^{-1} \text{ mol}^{-1} = 0.0820574587 \text{ L}\cdot\text{atm}\cdot\text{K}^{-1} \text{ mol}^{-1} = 0.083145 \text{ L}\cdot\text{bar}\cdot\text{K}^{-1}\cdot\text{mol}^{-1}$$

$$1 \text{ atm} = 101.325 \text{ kPa} = 1.01325 \text{ bar} = 14.696 \text{ psi} = 760 \text{ Torr} = 760 \text{ mmHg}$$

$$N_A = 6.022 \times 10^{23} \text{ mol}^{-1} \quad k = 8.62 \times 10^{-5} \text{ eV/K} \quad 1 \text{ eV} = 1.60 \times 10^{-19} \text{ J}$$

$$T(\text{K}) = T(\text{^{\circ}C}) + 273.15 \text{ K} \quad F = 9.6485309 \times 10^4 \text{ C mole}^{-1} \quad e = 1.60217733 \times 10^{-19} \text{ C}$$

$$\text{STP: } 273.15\text{K, 1 bar} \quad \text{SATP: } 298.15\text{K, 1 bar} \quad 1\text{L atm} = 101.325 \text{ J}$$

$$V = \frac{4\pi r^3}{3} \quad \rho = \frac{nA}{V_C N_A} \quad \%IC = \{1 - \exp[-0.25(X_A - X_B)^2]\} \times 100 \quad E = \int F dr \quad \frac{d}{dx}(x^n) = nx^{n-1}$$

$$APF = \frac{\text{TotalSphereVolume}}{\text{TotalUnitCellVolume}} \quad LD = \frac{\text{NumberOfAtomsCentredOnDirectionVector}}{\text{LengthOfDirectionVector}}$$

$$PD = \frac{\text{NumberOfAtomsCentredOnAPlane}}{\text{AreaOfPlane}} \quad DP = \frac{\overline{M_n}}{m} \quad \overline{M_n} = \sum x_i M_i \quad \overline{M_w} = \sum w_i M_i$$

$$N_V = N \exp(-\frac{Q_v}{kT}) \quad N_s = N \exp(-\frac{Q_s}{2kT}) \quad N_{fr} = N \exp(-\frac{Q_{fr}}{2kT})$$

$$E = \frac{\sigma}{\varepsilon} \quad \sigma = \frac{F}{A} \quad \epsilon = \frac{\Delta l}{l} \quad \tau = \frac{F}{A} \quad \tau = G\gamma \quad U_r = \frac{1}{2} \sigma_Y \epsilon_Y \quad \%CW = \left( \frac{A_0 - A_d}{A_0} \right) \times 100 \quad G = E/(2(1+\nu))$$

$$\sigma_y = \sigma_0 + k_y d^{-1/2} \quad \nu = -\frac{\epsilon_L}{\epsilon} = -\frac{\epsilon_x}{\epsilon_z} = -\frac{\epsilon_y}{\epsilon_z} \quad \tau_R = \sigma \cos\phi \cos\lambda \quad \sigma_y = \frac{\tau_{crss}}{(\cos\phi \cos\lambda)_{max}}$$

$$\theta = \cos^{-1} \left( \frac{u_1 u_2 + v_1 v_2 + w_1 w_2}{\sqrt{(u_1^2 + v_1^2 + w_1^2)(u_2^2 + v_2^2 + w_2^2)}} \right) \quad \sigma = n|e|\mu_e$$

$$V = IR \quad \rho = \frac{RA}{l} \quad \sigma = \frac{1}{\rho} \quad J = \sigma E \quad E = \frac{V}{l} \quad v_d = \mu_e E \quad \sigma = n|e|\mu_e + p|e|\mu_h$$

Idea gas equation of state:  $PV = nRT$

Van der Waals equation of state:

$$P = \frac{nRT}{(V - nb)} - \frac{an^2}{V^2}$$

First law, closed systems

$$\Delta U = q + w \Delta U = dq + dw$$

$$dw = -P_{\text{ext}} dV$$

$$dU = nC_{v,m}dT$$

$$H \equiv U + PV$$

$$dH = nC_{p,m}dT$$

For ideal gases,  $C_{p,m} = C_{v,m} + R$

Solids, Liquids,  $C_{p,m} = C_{v,m}$

$$G \equiv H - TS$$

$$dS \equiv \frac{dQ_{\text{reversible}}}{T}$$

For a process at constant temperature

$$\Delta G = \Delta H - T\Delta S$$

For an isothermal reversible process (ideal gas):

$$W_{\text{rev}} = - \int_{V_1}^{V_2} \frac{nRT}{V} dV = -nRT \ln \frac{V_2}{V_1} = -nRT \ln \frac{P_1}{P_2}$$

Adiabatic reversible process (ideal gas):

$$P_1 V_1^\gamma = P_2 V_2^\gamma \quad T_1 V_1^{(\gamma-1)} = T_2 V_2^{(\gamma-1)} \quad T_1 P_1^{[(1-\gamma)/\gamma]} = T_2 P_2^{[(1-\gamma)/\gamma]} \quad \left( \frac{\bar{C}_p}{\bar{C}_v} \right) = \gamma$$

$v_i$ : stoichiometric coefficient

Assuming no phase change, constant  $C_p$

$$\Delta H^\circ_{\text{rxn}} = \sum v_i * \Delta H^\circ_{f,i} + \Delta C_p * (T - 25^\circ\text{C})$$

$$\Delta C_p = \sum v_i * C_{p,i}$$

$$\text{Efficiency} = |w|/q_{in}$$

$$\epsilon(\text{ideal}) = 1 - T_C/T_H$$

For solids or liquids:

$$\text{Phase transition } \Delta S_{\text{trans}} = \frac{\Delta H_{\text{trans}}}{T_{\text{trans}}}$$

$$\Delta S^\circ(T_2) = \Delta S^\circ(T_1) + \int_{T_1}^{T_2} \Delta C_p \frac{dT}{T}$$

Standard entropy of the reaction  $\Delta S^\circ_{\text{rxn}} = \sum v_i * S_m^\circ$

Standard free energy of a reaction:  $\Delta G^\circ_{\text{rxn}} = \sum v_i * \Delta G^\circ_{f,i}$  or  $\Delta G^\circ_{\text{rxn}} = \Delta H^\circ_{\text{rxn}} - T\Delta S^\circ_{\text{rxn}}$

$$\Delta G^\circ = -RT \ln K \quad \ln \left( \frac{K_p(T_1)}{K_p(T_2)} \right) = -\frac{\Delta H^\circ_{\text{reaction}}}{R} \left( \frac{1}{T_1} - \frac{1}{T_2} \right) \quad (\text{assuming } \Delta H^\circ_{\text{reaction}} \text{ independent of } T)$$

Total free energy of the reaction  $aA + bB \Rightarrow cC + dD$

$$\Delta G_{\text{rxn}} = \Delta G_{\text{rxn}}^\emptyset + RT \ln(Q)$$

$$\text{where } Q = \left[ \frac{a_c^c a_d^d}{a_A^a a_B^b} \right] \text{ where } a = \text{activity} (\sim \text{concentration})$$

$$\Delta G_{\text{rxn}} = \Delta G_{\text{rxn}}^\emptyset + RT \ln(Q_p), \text{ where } Q_p = \left( \frac{P_C}{P^\circ} \right)^c \left( \frac{P_D}{P^\circ} \right)^d \left( \frac{P_A}{P^\circ} \right)^a \left( \frac{P_B}{P^\circ} \right)^b$$

$$\Delta G = -nFE \quad E = E^\circ - \frac{RT}{nF} \ln Q \quad E = E^\circ - \frac{0.0592}{n} \log Q \quad \text{at } 25^\circ\text{C} \quad I = \frac{nC}{t}$$

