

<i>Tutorial Section Number:</i>

FACULTY OF APPLIED SCIENCE AND ENGINEERING

FINAL EXAM

APRIL 2014

First Year

APS 104S

INTRODUCTION TO MATERIALS AND CHEMISTRY

Q1	/20
Q2	/20
Q3	/20
Q4	/20
Q5	/20
Q6	/20
Q7	/20
Q8	/20
Total	/140

Exam Type B

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

NAME: _____
Last First

STUDENT NO: _____

INSTRUCTIONS:

- This is a Type B examination. Only non-programmable calculators are allowed.
- *Answer any 7 of the 8 questions.*
- If all 8 questions are answered, only the first 7 will be marked.
- 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 where applicable clearly “box” your numerical results.
- 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*.

Name: _____

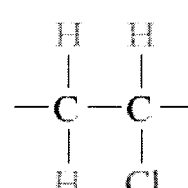
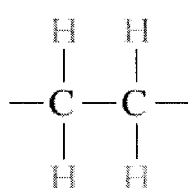
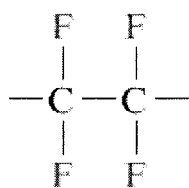
Student No: _____

QUESTION 1: (20 pts)

As appropriate circle the correct answer (from True/False or multiple choice), fill in the blank(s), provide a short answer with a sketch if required.

The following questions are worth one point each.

- (a) Geckos, harmless tropical lizards, have very sticky feet due to strong covalent bonding. **TRUE FALSE**
- (b) Ionic bonding is nondirectional. **TRUE FALSE**
- (c) The coordination number for FCC is 12 while the atomic packing factor is 0.74. **TRUE FALSE**
- (d) Circle all materials that form crystalline structures under normal solidification conditions:
- (i) Metals
 - (ii) Many ceramics
 - (iii) Certain polymers
 - (iv) All composites
 - (v) None of the above
- (e) Correctly label the following polymers [(polyethylene (PE), poly-vinyl-chloride (PVC), polytetrafluoroethylene (PTFE)]:



- (f) Circle the correct statement:
- (i) If $\Delta G > 0$ the process is exergonic;
 - (ii) If $\Delta G < 0$ the process is endergonic;
 - (iii) If $\Delta G = 0$ the process is exergonic and endergonic;
 - (iv) If $\Delta G < 0$ the process is exergonic;
 - (v) None of the above.
- (g) Ammonia, produced using the Haber process, is used extensively as a fertilizer. **TRUE FALSE**
- (h) Consider the following exothermic reaction: $\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightarrow 2\text{NH}_3(\text{g})$. If the reaction temperature is raised the reaction rate will _____.
- (i) Lithium ion batteries when being drained are an example of galvanic cells and when being charged an example of electrolytic cells. **TRUE FALSE**
- (j) The anode is
- (i) where oxidation occurs
 - (ii) where electrons are generated
 - (iii) the electrode with negative polarity
 - (iv) all of the above
 - (v) none of the above

Name: _____

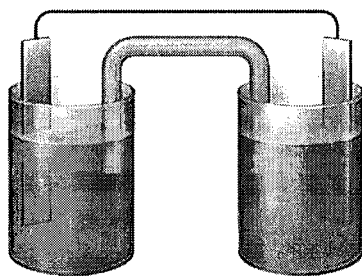
Student No: _____

The following questions are worth two points each.

- (k) Write the electronic configuration for aluminum:

- (l) The winter season this year has been the coldest that we have experienced in 25 years. One evidence of this is the large number of pot holes in the streets of Toronto. Explain why this is the case (describing the properties of the relevant material(s) and the processes thereof).

- (m) Consider the concentration cell based on the Cd/Cd^{2+} reaction shown below. In which direction will the electrons flow (*left to right* OR *right to left*)? What will be the cell voltage at equilibrium?



☐ *left to right*

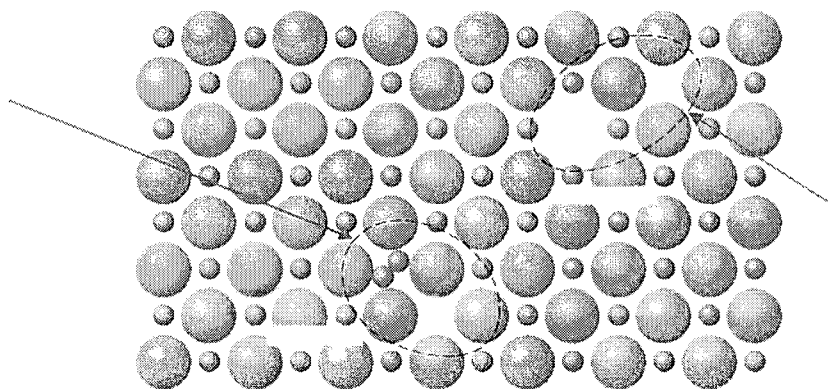
☐ *right to left*

$E_{\text{cell} - \text{equilibrium}} =$ _____

Cd/Cd^{2+} (0.1 M)

Cd/Cd^{2+} (1.0 M)

- (n) Identify the two defects shown in the diagram below.



- (o) Consider the reaction $\text{H}_2 + \frac{1}{2}\text{O}_2 \rightarrow \text{H}_2\text{O}$ with an enthalpy of reaction of 286 kJ/mol and Gibbs free energy of reaction of 237 kJ/mol.
- (i) The reaction is carried out in a combustion chamber with $T_h = 950$ K and $T_c = 400$ K. Calculate the maximum thermal work that can be extracted.
- (ii) The reaction is carried out in an electrolytic fuel cell. Calculate the maximum electrical work that can be extracted.

Name: _____

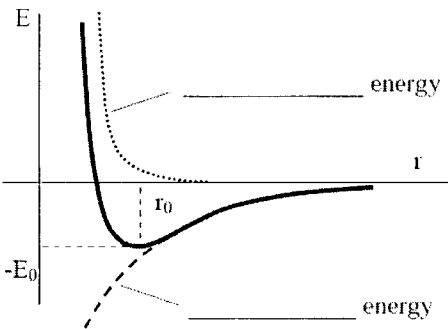
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QUESTION 2: (20 pts)

A. State the type of bonding (ionic, covalent, metallic, van der Waals, hydrogen-bonding) for each of the materials listed in the table below. *Note:* Look at both the bond energy and substance. (3 marks)

Substance	Bond Energy (eV/(atom,ion,molecule))	Bonding Type
H ₂ O	0.52	
NaCl	3.3	
Si	4.7	
Ar	0.08	
Fe	4.2	
NH ₃	0.36	

B. The figure on the right shows the total potential energy curve (in bold line) which represents the interaction between two atoms as a function of the inter-atomic separation. The total potential energy curve is a sum of two different two types of potential energy interactions shown as dashed and dotted curves. (2 marks)



- i) Label the two types of potential energy on the figure;
- ii) State the equilibrium interatomic distance: _____;
- iii) State the binding energy between the atoms: _____.

C. The potential energy between Na⁺ and Cl⁻ ions in a NaCl crystal can be written as:

$$E(r) = -\frac{A}{r} + \frac{B}{r^8}$$

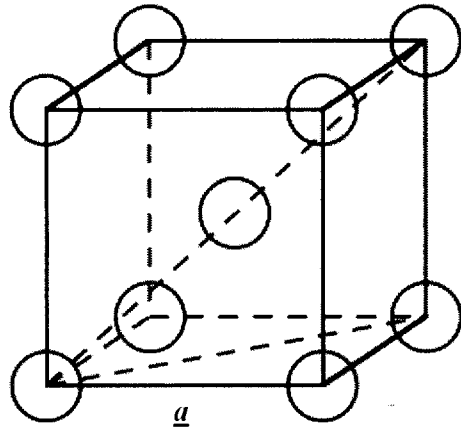
where $A = 4.03 \times 10^{-28} \text{ J}\cdot\text{m}$ and $B = 6.97 \times 10^{-96} \text{ J}\cdot\text{m}^8$. The energy is given in Joules per ion pair. Calculate the equilibrium interatomic separation between the ions. (5 marks)

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D.

- (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 planar atomic density in the (110) plane of the BCC crystal structure. [3 marks]

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QUESTION 3: (20 pts)

A..Fill-in the blanks (1 mark each):

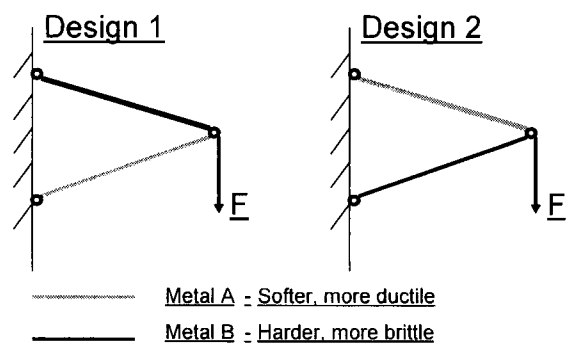
- a. The reversible, non permanent deformation is called _____ deformation.
- b. For a material with poisson's ratio of 0.5, it's volume change is _____.
- c. Ductility is a measure of the degree of _____ deformation that has been sustained at fracture.
- d. In general, the elastic modulus of ceramics is _____ than that of polymers. Ceramics are dominated by _____ bonds, while polymers are dominated by _____ bonds.
- e. For a material with a poisson's ratio of 0.3, its shear modulus G is about _____ E .

B..Answer the following questions:

- (i) Two cylindrical nickel wires A and B have diameter ratio of 1:2 (i.e. $d_A:d_B = 1:2$). What is the ratio of their strain ($\epsilon_A : \epsilon_B$) when the same force is applied along the long axis? Assume that the deformation is elastic. **(2 marks)**

- (ii) State two reasons where porosity negatively influences the flexural strength of ceramic materials. **(2 marks)**

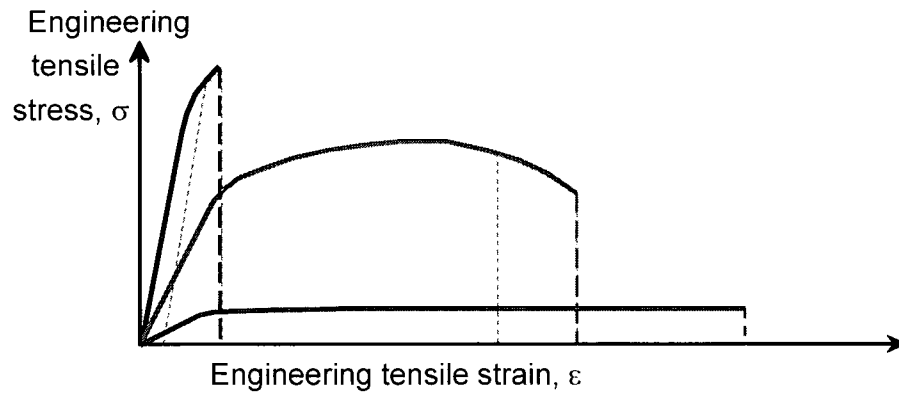
- (iii) The figure on the right shows two designs with the same parameters. Which one is better and why? **(2 marks)**



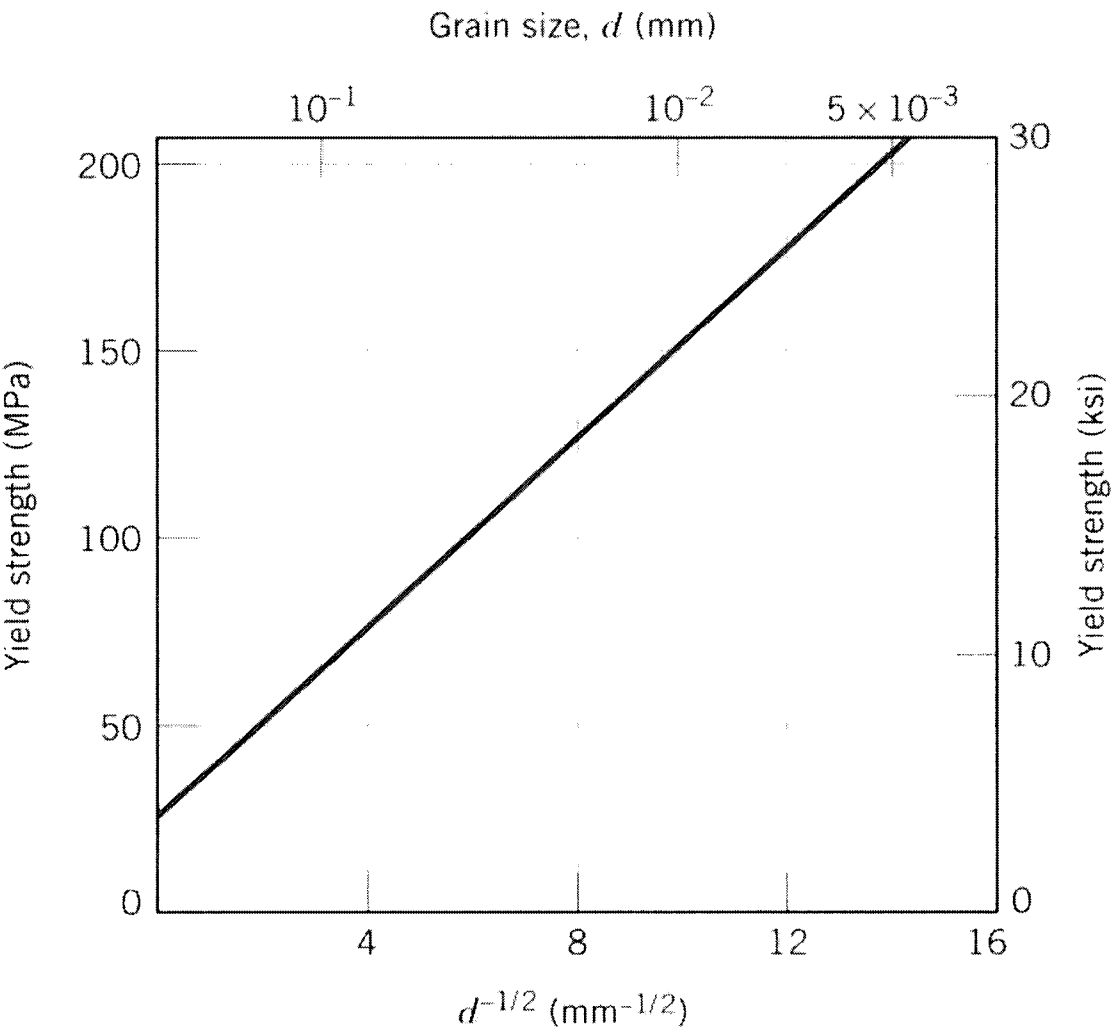
Name: _____

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C...With reference to the following stress-strain curves:



- (i) Identify which curve represents the following materials: ceramic, metal and polymer. Clearly indicate the material on the graph. **(3 marks)**
 - (ii) Mark the tensile strength on the stress-strain curve that represents metals. **(1 mark)**
- D.** Describe the following strengthening mechanisms, including an explanation as to how dislocations are involved in each of the strengthening techniques. **(3 marks - 1 mark each)**
- (i) Grain size reduction
 - (ii) Solid-solution strengthening
 - (iii) Strain hardening
- E.** From the plot below (see next page) of yield strength versus $(\text{grain diameter})^{-1/2}$ for a 70Cu–30Zn cartridge brass, determine the values for the constants σ_0 and k_y in the equation $\sigma_y = \sigma_0 + k_y d^{-1/2}$ **(3 marks)**



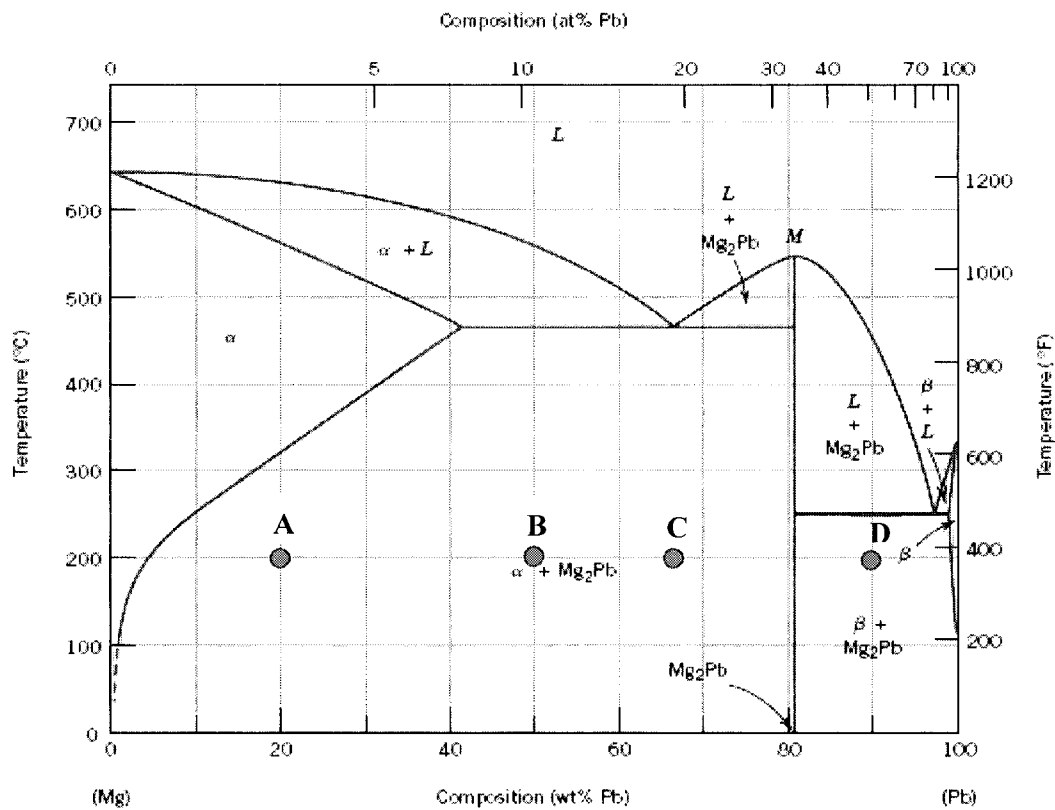
Plot showing the influence of grain size on the yield strength of a 70Cu-30Zn brass alloy. Note that the grain diameter increases from right to left and is not linear.

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QUESTION 4: (20 pts)

A. The magnesium-lead diagram is shown below. Use it to answer questions (a) – (e).



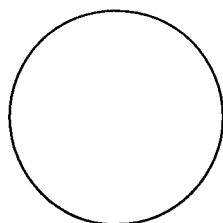
- (a) Label clearly (i) the two eutectic points and (ii) the congruent point on the diagram above. [3 pts]
(b) What are the temperatures at the two eutectic isotherms? [2 pts]

- (c) Write the eutectic reaction for the eutectic transformation that is occurring at
(i) the higher eutectic isotherm, and
(ii) the lower eutectic isotherm.

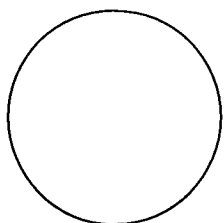
Specify the composition of all the phases. [4 pts]

- (d) At the eutectic composition (associated with $T_{E(\text{higher})}$), determine the amounts of each phase present, just below the eutectic isotherm. [2 pts]

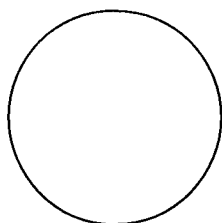
- (e) For points A-D on the graph above, sketch the correspond microstructures in the circles provided below. [4 pts]



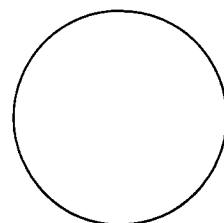
A



B



C



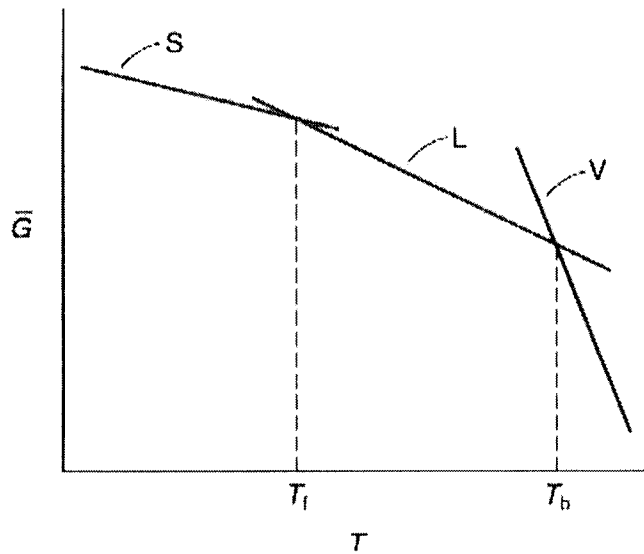
D

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B.

- (f) **True or False?** Consider water and ice at equilibrium at some T and P , it then follows that $G_{\text{ice}} = G_{\text{water}}$. [1 pts]
- (g) **True or False?** The chemical potential of water is higher than the chemical potential of an aqueous glucose solution. [1 pts]
- (h) On the diagram below, if the pressure is increased, sketch the effect that it would have on the boiling point of this substance (i.e., sketch the new curves). What is the change in the boiling point? [3 pts]



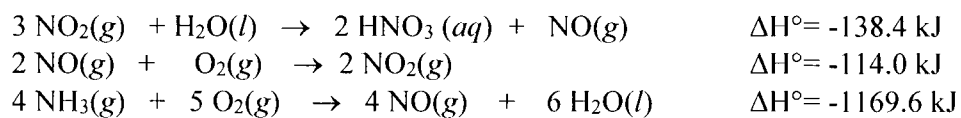
Name: _____

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QUESTION 5: (20 pts)

(a) Write down the ΔH°_f reaction for $\text{HNO}_3(aq)$ [that is, chemical reaction for the standard enthalpy of formation of $\text{HNO}_3(aq)$]. [2 pts]

(b) Set up a Hess's law cycle, and use the following information to calculate the ΔH°_f for aqueous nitric acid, $\text{HNO}_3(aq)$. You will need to use fractional coefficient for some equations. [6 pts]



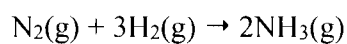
Given: $\text{NH}_3(g)$ $\Delta H^\circ_f = -46.1 \text{ kJ/mol}$
 $\text{H}_2\text{O}(l)$ $\Delta H^\circ_f = -285.8 \text{ kJ/mol}$

(c) Calculate ΔU for the reaction in (a). If you were not able to solve part (b) use $\Delta H^\circ_f = -200 \text{ kJ/mol}$. [2 pts]

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(d) Use $\text{NH}_3(\text{g})$ $\Delta H^\circ_f = -46.1 \text{ kJ/mol}$ to calculate ΔH of the following reaction at 1000 K



with $C_{p,m} = 3.502R$, $3.466R$, and $4.217R$ for $\text{N}_2(\text{g})$, $\text{H}_2(\text{g})$, and $\text{NH}_3(\text{g})$, respectively. [6 pts]

- (e) In a calorimetry experiment, 5.00 g of ammonia ($\Delta H^\circ_f = -46.1 \text{ kJ/mol}$) are produced after stoichiometric amounts of nitrogen and hydrogen has reacted. Determine the combined heat capacity of the calorimeter and water, if the calorimeter and water-bath temperature rises by 0.42°C as a result of the production of 5.00 g of ammonia. [4 pts]

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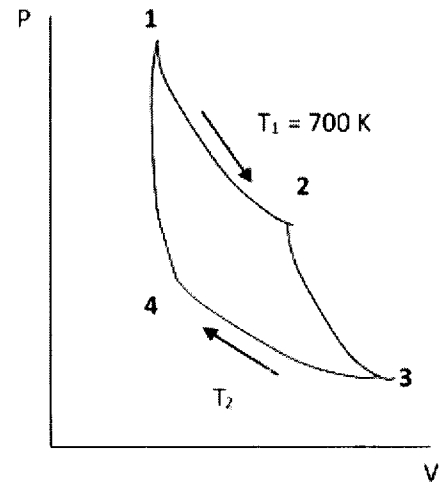
Student No: _____

QUESTION 6: Engle-Reid Chapter 5 (20 pts)

Consider the Carnot cycle shown on the right for 0.6 moles of an ideal *monatomic* gas for which you are given:

$$T_1 = 700 \text{ K}, V_2 = 2V_1 \text{ and } V_3 = 1.5V_2$$

(a) Calculate $w_{1 \rightarrow 2}$, $\Delta U_{1 \rightarrow 2}$, $q_{1 \rightarrow 2}$. [4 pts]



(b) Find T_2 . [2 pts]

(c) What is the efficiency of a Carnot engine operating on this cycle? [2 pts]

(d) What is the total work around the cycle? [2 pts]

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(e) How much heat is rejected at T_2 ? [2 pts]

(f) Evaluate $\Delta S_{3 \rightarrow 4}$. [2 pts]

(g) Describe on the molecular scale the sign of $\Delta S_{3 \rightarrow 4}$ calculated in (f). [3 pts]

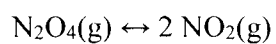
(h) What is $\Delta S_{\text{universe}}$ for the whole cycle? Show your calculation or explain your result. [3 pts]

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QUESTION 7: (20 pts)

(A) Dinitrogen tetroxide is 18.46 per cent dissociated at 25°C and 1.00 bar in the equilibrium



Calculate/determine the following:

(i) K at 25°C [6 pts]

(ii) $\Delta_r G^\ominus$ [3 pts]

(iii) $\Delta_r G$ [2 pts]

(iv) K at 100°C given that $\Delta_r H^\ominus = +57.2 \text{ kJ mol}^{-1}$ over the temperature range [3 pts]

Name: _____ Student No: _____

(B) The standard Gibbs energy of formation of $\text{NH}_3(\text{g})$ is $-16.5 \text{ kJ mol}^{-1}$ at 298 K.

- (i) What is the reaction Gibbs energy when the partial pressure of the N_2 , H_2 , and NH_3 (treated as ideal gases) are 3.0 bar, 1.0 bar, and 4.0 bar, respectively? **[3 pts]**

- (ii) What is the spontaneous direction of the reaction in this case? Explain your answer. **[3 pts]**

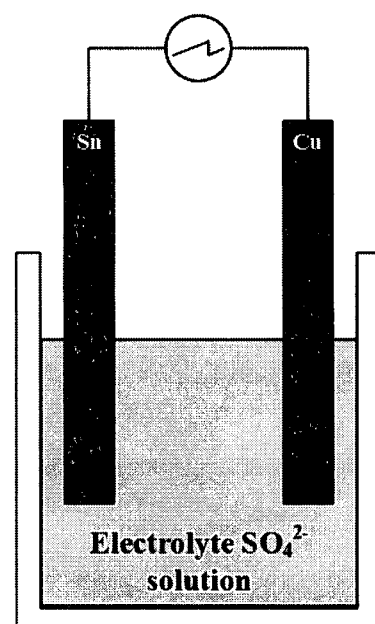
Name: _____

Student No: _____

QUESTION 8: (20 pts)

(A) Consider the electrochemical cell on the right and the following data (at 25 °C):

	E^\ominus
$\text{Cu}^{2+} + 2\text{e}^- \rightarrow \text{Cu(s)}$	+0.34 V
$\text{Sn}^{2+} + 2\text{e}^- \rightarrow \text{Sn(s)}$	-0.14 V



(i) Assuming that the electrolyte solution could be **either** CuSO_4 or SnSO_4 , what is the spontaneous/voltaic electrochemical reaction that can occur for this cell? [4 pts]

(ii) What should be the electrolyte for the voltaic cell? [1 pt]

(iii) Which electrode is the cathode for the voltaic cell? [1 pt]

(iv) Which electrode is the anode for the voltaic cell? [1 pt]

(v) What is the equilibrium constant for the spontaneous reaction at 25 °C? [4 pts]

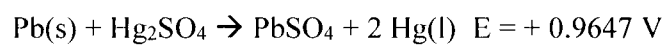
(vi) What will physically happen to the cathode during this process? [1 pt]

(vii) What will physically happen to the anode during this process? [1 pt]

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(B) Consider the following electrochemical reaction:



And given $\left(\frac{\partial E}{\partial T}\right)_p = 1.74 \times 10^{-4} \text{ V K}^{-1}$

(i) Calculate ΔG_r for this reaction. **[2 pts]**

(ii) Calculate ΔS_r for this reaction. **[2 pts]**

(iii) Calculate ΔH_r for this reaction. **[2 pts]**

(iv) Is this process enthalpically or entropically driven? Why? **[1 pts]**

FORMULAE & CONSTANTS (You may tear these sheets off.)

$R = 8.3145 \text{ J}\cdot\text{K}^{-1}\cdot\text{mol}^{-1} = 0.0820574587 \text{ L}\cdot\text{atm}\cdot\text{K}^{-1}\cdot\text{mol}^{-1} = 0.083145 \text{ L}\cdot\text{bar}\cdot\text{K}^{-1}\cdot\text{mol}^{-1}$ $1 \text{ atm} = 101.325 \text{ kPa} = 1.01325 \text{ bar} = 14.696 \text{ psi} = 760 \text{ Torr} = 760 \text{ mmHg}$ $N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$ $k = 8.62 \times 10^{-5} \text{ eV/K}$ $1 \text{ eV} = 1.60 \times 10^{-19} \text{ J}$ $T(\text{K}) = T(^{\circ}\text{C}) + 273.15 \text{ K}$ $F = 9.6485309 \times 10^4 \text{ C mole}^{-1}$ $e = 1.60217733 \times 10^{-19} \text{ C}$ STP: 273.15K, 1 atm SATP: 298.15K, 1 bar 1L atm = 101.325 J	
$V = \frac{4\pi r^3}{3}$ $\rho = \frac{nA}{V_C N_A}$ $\%IC = \{1 - \exp[-0.25(X_A - X_B)^2]\} \times 100$ $E = \int F dr$ $\frac{d}{dx}(x^n) = nx^{n-1}$ $APF = \frac{\text{Total Sphere Volume}}{\text{Total Unit Cell Volume}}$ $LD = \frac{\text{Number Of Atoms Centred On Direction Vector}}{\text{Length Of Direction Vector}}$ $PD = \frac{\text{Number Of Atoms Centred On A Plane}}{\text{Area Of Plane}}$ $DP = \frac{\overline{M_n}}{m}$ $\overline{M_n} = \sum x_i M_i$ $\overline{M_w} = \sum w_i M_i$ $N_V = N \exp(-\frac{Q_v}{kT})$ $N_S = N \exp(-\frac{Q_s}{2kT})$ $N_{fr} = N \exp(-\frac{Q_{fr}}{2kT})$ $E = \frac{\sigma}{\epsilon}$ $\sigma = \frac{F}{A}$ $\epsilon = \frac{\Delta l}{l}$ $\tau = \frac{F}{A}$ $\tau = G\gamma$ $U_r = \frac{1}{2} \sigma_Y \epsilon_Y$ $\%CW = \left(\frac{A_0 - A_d}{A_0}\right) \times 100$ $G = E/(2(1 + \nu))$ $\sigma_y = \sigma_0 + k_y d^{-1/2}$ $\nu = -\frac{\epsilon_L}{\epsilon} = -\frac{\epsilon_x}{\epsilon_z} = -\frac{\epsilon_y}{\epsilon_z}$ $\tau_R = \sigma \cos\phi \cos\lambda$ $\sigma_y = \frac{\tau_{crss}}{(\cos\phi \cos\lambda)_{max}}$ $\theta = \cos^{-1} \left(\frac{u_1 u_2 + v_1 v_2 + w_1 w_2}{\sqrt{(u_1^2 + v_1^2 + w_1^2)(u_2^2 + v_2^2 + w_2^2)}} \right)$ $\sigma = n e \mu_e$ $V = IR$ $\rho = \frac{RA}{l}$ $\sigma = \frac{1}{\rho}$ $J = \sigma E$ $E = \frac{V}{l}$ $v_d = \mu_e E$ $\sigma = n e \mu_e + p e \mu_h$	
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 $\Delta U = q + w$ $dU = dq + dw$ $dw = -P_{ext}dV$ $dU = nC_{v,m}dT$	$H \equiv U + PV$ $dH = nC_{p,m}dT$ For ideal gases, $C_{p,m} = C_{v,m} + R$ Solids, Liquids, $C_{p,m} = C_{v,m}$ $G \equiv H - TS$ $dS \equiv \frac{dQ_{reversible}}{T}$ For a process at constant temperature $\Delta G = \Delta H - T\Delta S$
For an isothermal reversible process (ideal gas): $W_{rev} = -\int_{V_i}^{V_f} \frac{nRT}{V} dV = -nRT \ln \frac{V_f}{V_i} = -nRT \ln \frac{P_i}{P_f}$ Adiabatic reversible process (ideal gas): $P_1 V_1^\gamma = P_2 V_2^\gamma$ $T_1 V_1^{(\gamma-1)} = T_2 V_2^{(\gamma-1)}$ $T_1 P_1^{[(1-\gamma)/\gamma]} = T_2 P_2^{[(1-\gamma)/\gamma]}$ $\left(\frac{\overline{C_p}}{\overline{C_v}}\right) = \gamma$	ν_i : stoichiometric coefficient Assuming no phase change, constant C_p $\Delta H^{\circ}_{rxn} = \sum \nu_i \Delta H^{\circ}_{f,i} + \Delta C_p (T - 25^{\circ}\text{C})$ $\Delta C_p = \sum \nu_i C_{p,i}$ Efficiency = $ w /q_{in}$ $\epsilon(\text{ideal}) = 1 - T_C/T_H$
For solids or liquids: Phase transition $\Delta S_{trans} = \frac{\Delta H_{trans}}{T_{trans}}$ $\Delta S^{\circ}(T_2) = \Delta S^{\circ}(T_1) + \int_{T_1}^{T_2} \Delta C_p \frac{dT}{T}$ Standard entropy of the reaction $\Delta S^{\circ}_{rxn} = \sum \nu_i S_{m,i}^{\circ}$ Standard free energy of a reaction: $\Delta G^{\circ}_{rxn} = \sum \nu_i \Delta G^{\circ}_{f,i}$ or $\Delta G^{\circ}_{rxn} = \Delta H^{\circ}_{rxn} - T\Delta S^{\circ}_{rxn}$ $\Delta G^{\circ} = -RT \ln K$ $\ln \left(\frac{K_p(T_1)}{K_p(T_2)} \right) = -\frac{\Delta H^{\circ}_{reaction}}{R} \left(\frac{1}{T_1} - \frac{1}{T_2} \right)$ (assuming $\Delta H^{\circ}_{reaction}$ independent of T)	$\Delta S = nR \ln \frac{V_f}{V_i}$ (isothermal) $\Delta S = nC_v \ln \frac{T_f}{T_i}$ (change in T at const V) $\Delta S = nC_p \ln \frac{T_f}{T_i}$ (change in T at const P)
Total free energy of the reaction $aA + bB \Rightarrow cC + dD$ $\Delta G_{rxn} = \Delta G_{rxn}^{\circ} + RT \ln(Q)$ where $Q = \left[\frac{a_c^c a_d^d}{a_A^a a_B^b} \right]$ where a = activity (~ concentration)	$\Delta G_{rxn} = \Delta G_{rxn}^{\circ} + RT \ln(Q_p)$, where $Q_p = \frac{\left(\frac{P_C}{P^{\circ}}\right)^c \left(\frac{P_D}{P^{\circ}}\right)^d}{\left(\frac{P_A}{P^{\circ}}\right)^a \left(\frac{P_B}{P^{\circ}}\right)^b}$
$\Delta G = -nFE$ $E = E^{\circ} - \frac{RT}{nF} \ln Q$ $E = E^{\circ} - \frac{0.0592}{n} \log Q$ at 25°C $l = \frac{nC}{t}$	
$\Delta S_r = nF \left(\frac{\partial E}{\partial T} \right)_p$ $K = \frac{[C]^c [D]^d}{[A]^a [B]^b}$ $\ln K = \frac{-\Delta G^{\circ}}{RT}$ $\ln K = \frac{nFE^{\circ}}{RT}$	

PERIODIC TABLE OF THE ELEMENTS

<http://www.ktf-split.hr/periodni/en/>

PERIOD	GROUP																18 VIIIA	
	1 IA	2 IIA											13 IIIA	14 IVA	15 VA	16 VIA	17 VIIA	18 VIIIA
1	1 1.0079 H HYDROGEN																	2 4.0026 He HELIUM
2	3 6.941 Li LITHIUM	4 9.0122 Be BERYLLIUM											5 10.811 B BORON	6 12.011 C CARBON	7 14.007 N NITROGEN	8 15.999 O OXYGEN	9 18.998 F FLUORINE	10 20.180 Ne NEON
3	11 22.990 Na SODIUM	12 24.305 Mg MAGNESIUM											13 26.982 Al ALUMINIUM	14 28.086 Si SILICON	15 30.974 P PHOSPHORUS	16 32.065 S SULPHUR	17 35.453 Cl CHLORINE	18 39.948 Ar ARGON
4	19 39.098 K POTASSIUM	20 40.078 Ca CALCIUM	21 44.956 Sc SCANDIUM	22 47.867 Ti TITANIUM	23 50.942 V VANADIUM	24 51.996 Cr CHROMIUM	25 54.938 Mn MANGANESE	26 55.845 Fe IRON	27 58.933 Co COBALT	28 58.693 Ni NICKEL	29 63.546 Cu COPPER	30 65.39 Zn ZINC	31 69.723 Ga GALLIUM	32 72.64 Ge GERMANIUM	33 74.922 As ARSENIC	34 78.96 Se SELENIUM	35 79.904 Br BROMINE	36 83.80 Kr KRYPTON
5	37 85.468 Rb RUBIDIUM	38 87.62 Sr STRONTIUM	39 88.906 Y YTTRIUM	40 91.224 Zr ZIRCONIUM	41 92.906 Nb NIOBIUM	42 95.94 Mo MOLYBDENUM	43 (98) Tc TECHNETIUM	44 101.07 Ru RUTHENIUM	45 102.91 Rh RHODIUM	46 106.42 Pd PALLADIUM	47 107.87 Ag SILVER	48 112.41 Cd CADMIUM	49 114.82 In INDIUM	50 118.71 Sn TIN	51 121.76 Sb ANTIMONY	52 127.60 Te TELLURIUM	53 126.90 I IODINE	54 131.29 Xe XENON
6	55 132.91 Cs CAESIUM	56 137.33 Ba BARIUM	57-71 La-Lu Lanthanide	72 178.49 Hf HAFNIUM	73 180.95 Ta TANTALUM	74 183.84 W TUNGSTEN	75 186.21 Re RHENIUM	76 190.23 Os OSMIUM	77 192.22 Ir IRIDIUM	78 195.08 Pt PLATINUM	79 196.97 Au GOLD	80 200.59 Hg MERCURY	81 204.38 Tl THALLIUM	82 207.2 Pb LEAD	83 208.98 Bi BISMUTH	84 (209) Po POLONIUM	85 (210) At ASTATINE	86 (222) Rn RADON
7	87 (223) Fr FRANCIUM	88 (226) Ra RADIUM	89-103 Ac-Lr Actinide	104 (261) Rf RUTHERFORDIUM	105 (262) Db DUBNIUM	106 (266) Sg SEABORGIUM	107 (264) Bh BOHRIUM	108 (277) Hs HASSIUM	109 (268) Mt MEITNERIUM	110 (281) Uun UNUNNIUM	111 (272) Uuu UNUNUNIUM	112 (285) Uub UNUBIUM		114 (289) Uuq UNUNQUADIUM				

LANTHANIDE

57 138.91 La LANTHANUM	58 140.12 Ce CERIUM	59 140.91 Pr PRASEODYMIUM	60 144.24 Nd NEODYMIUM	61 (145) Pm PROMETHIUM	62 150.36 Sm SAMARIUM	63 151.96 Eu EUROPIUM	64 157.25 Gd GADOLINIUM	65 158.93 Tb TERBIUM	66 162.50 Dy DYSPROSIUM	67 164.93 Ho HOLMIUM	68 167.26 Er ERBIUM	69 168.93 Tm THULIUM	70 173.04 Yb YTTERBIUM	71 174.97 Lu LUTETIUM
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ACTINIDE

89 (227) Ac ACTINIUM	90 232.04 Th THORIUM	91 231.04 Pa PROTACTINIUM	92 238.03 U URANIUM	93 (237) Np NEPTUNIUM	94 (244) Pu PLUTONIUM	95 (243) Am AMERICIUM	96 (247) Cm CURIUM	97 (247) Bk BERKELIUM	98 (251) Cf CALIFORNIUM	99 (252) Es EINSTEINIUM	100 (257) Fm FERMIUM	101 (258) Md MENDELEVIUM	102 (259) No NOBELIUM	103 (262) Lr LAWRENCIUM
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(1) Pure Appl. Chem., 73, No. 4, 667-683 (2001)

Relative atomic mass is shown with five significant figures. For elements having no stable nuclides, the value enclosed in brackets indicates the mass number of the longest-lived isotope of the element.

However three such elements (Th, Pa, and U) do have a characteristic terrestrial isotopic composition, and for these an atomic weight is tabulated.

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