Exam3Fall2009thermoelectro

Multiple Choice

Identify the letter of the choice that best completes the statement or answers the question.

- 1. Thermodynamics can be used to determine all of the following EXCEPT
 - a. the direction in which a reaction is spontaneous.
 - b. the extent to which a reaction occurs.
 - c. the rate of reaction.
 - d. the temperature at which a reaction is spontaneous.
 - e. the enthalpy change of a reaction.
- 2. Which of the following involves a decrease in entropy?
 - a. the sublimation of carbon dioxide
 - b. the dissolution of NaCl in water
 - c. the decomposition of $N_2O_4(g)$ to $NO_2(g)$
 - d. the evaporation of ethanol
 - e. the freezing of liquid water into ice
- 3. Of the following product-favored processes, which are endothermic?
 - 1. the combustion of methane to produce water and carbon dioxide
 - 2. the expansion of an ideal gas
 - 3. the melting of ice at temperatures greater than 0°C.
 - a. 1 only
 - b. 2 only
 - c. 3 only
 - d. 1 and 2
 - e. 2 and 3
- 4. Calculate the standard molar entropy change for the combustion of methane.

$$CH_4(g) + 2 O_2(g) \rightarrow CO_2(g) + 2 H_2O(g)$$

Species	S^{o} (J/K·mol)
CH ₄ (g)	186.3
$O_2(g)$	205.1
$CO_2(g)$	213.7
$H_2O(g)$	188.8

- a. -5.2 J/K
- b. -1.0 J/K
- c. +1.0 J/K
- d. +5.2 J/K
- e. +11.1 J/K
- 5. Predict the signs of ΔH , ΔS , and ΔG for the melting of ice at 50°C.
 - a. $\Delta H < 0, \Delta S < 0, \Delta G < 0$
 - b. $\Delta H < 0, \Delta S > 0, \Delta G < 0$
 - c. $\Delta H < 0, \Delta S > 0, \Delta G < 0$
 - d. $\Delta H > 0$, $\Delta S < 0$, $\Delta G < 0$
 - e. $\Delta H > 0$, $\Delta S > 0$, $\Delta G < 0$

 6.	Above what tem	nperature would you	expect a reaction to become spontaneous if $\Delta H = +322$ kJ and $\Delta S = +531$
	J/K?		
	a. 171 Kb. 209 K		
	c. 606 K		
		n will be spontaneou	s at any temperature.
			aneous at any temperature.
7.	Calculate ΔG_{mm}^*	for the reaction belo	ow at 25.0°C
	IXI		
	$2 H_2S(g) + O_2(g)$	$(x) \rightarrow 2 H_2O(g) + S(s)$	
	given $\Delta H_{\text{man}}^* = -4$	442.4 kJ, and $\Delta S_{\mathtt{man}}^*$ =	= -175.4 J/K.
	a438.0 kJ		
	b390.1 kJ		
	c321.9 kJ		
	d. +3943 kJ e. +5182 kJ		
0		C .1 1.1	. 25 000
 8.	Calculate AG	for the reaction belo	w at 25.0°C.
	$4 \text{ Fe(s)} + 3 \text{ O}_2(\text{g})$	$(s) \rightarrow 2 \text{ Fe}_2\text{O}_3(s)$	
	1 (1)	2 - 3(-)	
	Species	$\Delta H_f^*(\mathrm{kJ/mol})$	$S_f^*(J/K \cdot mol)$
	Fe(s)	0	27.78
	$O_2(g)$	0	205.14
	$Fe_2O_3(s)$	-824.2	87.40
	a1629 kJ		
	b1484 kJ		
	c780.8 kJ		
	d659.7 kJ		
	e. +1629 kJ		
 9.	The free energy	change for the form	ation of the complex ion AlF ₆ ³ - is -140. kJ at 25°C. What is the
		stant for the reaction	?
	a. 2.9×10^{-25}		
	b. 5.65×10^{1}		
	c. 3.5×10^{24} d. 5.2×10^{29}		
	e. 2.3×10^{56}		
10.	In the following	roation	
 10.	in the following	, reaction,	
	$Fe^{2+}(aq) + Ag^{+}(a$	$aq) \rightarrow Fe^{3+}(aq) + Ag$	$g(\mathbf{s})$
	a. Fe^{2+} is oxidi	ized and Fe ³⁺ is redu	ced.
		ized and Ag ⁺ is reduce	
		zed and $Ag(s)$ is red	
	d. Ag^{+} is oxidi	zed and Fe ²⁺ is reduce	ced.
	_	zed and Fe ³⁺ is redu	
 11.	Write a balance	d chemical equation	for the following reaction in a basic solution.

$$\begin{aligned} &H_2O_2(aq) + Cr(OH)_3(s) \rightarrow H_2O(\ell) + CrO_4^{-2}(aq) \\ &a. & 2 H_2O_2(aq) + 3 Cr(OH)_3(s) \rightarrow H_2O(\ell) + 3 CrO_4^{-2}(aq) + 11/2 H'(aq) \\ &b. & 2 H_2O_3(aq) + Cr(OH)_3(s) \rightarrow H_2O(\ell) + CrO_4^{-2}(aq) + 2 OH(aq) \\ &c. & H_2O_3(aq) + 2 Cr(OH)_3(s) \rightarrow H_2O(\ell) + 2 CrO_4^{-2}(aq) + 4 H_2O(\ell) \\ &d. & 3 H_2O_2(aq) + 2 Cr(OH)_3(s) \rightarrow 2 H_2O(\ell) + 2 CrO_4^{-2}(aq) + 8 H_2O(\ell) \\ &e. & 4 H_2O_3(aq) + 2 Cr(OH)_3(s) \rightarrow 2 H_2O(\ell) + 2 CrO_4^{-2}(aq) + 4 OH'(aq) \\ &12. & \text{All of the following statements concerning voltaic cells are true EXCEPT} \\ &a. & \text{the two half-cells are connected by a salt bridge.} \\ &b. & \text{electrons flow from the anode to the cathode.} \\ &c. & \text{oxidation occurs at the cathode.} \\ &d. & \text{voltaic cells can be used as a source of energy.} \\ &e. & \text{a voltatic eell consists of two-half cells.} \\ &13. & \text{What is the correct cell notation for the reaction below?} \\ &Cu^{2+}(aq) + Pb(s) \rightarrow Cu(s) + Pb^{2+}(aq) \\ &a. & Pb \mid Pb^{2+}(aq) \mid Cu^{2+}(aq) \mid Cu \\ &b. & Pb \mid Cu^{2+}(aq) \mid Pb^{2+}(aq) \mid Cu^{2+} \\ &d. & Cu \mid Pb^{2+}(aq) \mid Hb^{2+}(aq) \mid Cu^{2+} \\ &d. & Cu \mid Pb^{2+}(aq) \mid Hb^{2+}(aq) \mid Db \\ &e. & Cu \mid Cu^{2+}(aq) \mid Bb^{2+}(aq) \mid Cd^{2+}(aq) \mid Pb \\ &e. & Cu \mid Cu^{2+}(aq) \mid Sh^{4+}(aq) \rightarrow Cd(s) + Sh^{4+}(aq) \\ &b. & Cd^{2+}(aq) + Sh^{2+}(aq) \rightarrow Cd(s) + Sh^{4+}(aq) \\ &b. & Cd^{2+}(aq) + Sh^{2+}(aq) \rightarrow Cd(s) + Sh^{4+}(aq) \\ &b. & Cd^{2+}(aq) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq) \rightarrow Cd^{2+}(aq) + Sh^{4+}(aq) \\ &c. & Cd(s) + Sh^{2+}(aq)$$

____ 16. Use the standard reduction potentials below to determine which compound or ion is the best reducing agent?

Hg²⁺(aq) + 2 e⁻ → Hg(
$$\ell$$
) E° = +0.855 V
Cu²⁺(aq) + 2 e⁻ → Cu(s) E° = +0.337 V
Cd²⁺(aq) + 2 e⁻ → Cd(s) E° = -0.40 V

a. Hg^{2+}

- b. $Hg(\ell)$
- c. Cu²⁺
- d. Cd^{2+}
- e. Cd
- 17. Consider the following half-reactions:

Fe³⁺(aq) + e⁻
$$\rightarrow$$
 Fe²⁺(aq) $E^{\circ} = +0.77 \text{ V}$
Sn²⁺(aq) + 2 e⁻ \rightarrow Sn(s) $E^{\circ} = -0.14 \text{ V}$
Fe²⁺(aq) + 2 e⁻ \rightarrow Fe(s) $E^{\circ} = -0.44 \text{ V}$
Al³⁺(aq) + 3 e⁻ \rightarrow Al(s) $E^{\circ} = -1.66 \text{ V}$
Mg²⁺(aq) + 2 e⁻ \rightarrow Mg(s) $E^{\circ} = -2.37 \text{ V}$

Which of the above metals or metal ions are able to oxidize Al(s)?

- a. Fe^{3+} and Sn^{2+}
- b. Fe³⁺, Sn²⁺, and Fe²⁺
 c. Fe²⁺, Sn, and Fe
- d. Mg and Mg²⁺
- e. Mg^{2+} only
- 18. Given the following two half-reactions, determine which overall reaction is spontaneous and calculate the cell potential.

$$Mg^{2+}(aq) + 2 e^{-} \rightarrow Mg(s)$$
 $E^{\circ} = -2.37 \text{ V}$
Ni²⁺(aq) + 2 e⁻ \rightarrow Ni(s) $E^{\circ} = -0.25 \text{ V}$

a.
$$Mg^{2+}(aq) + Ni(s) \rightarrow Mg(s) + Ni^{2+}(aq)$$
 $E_{cell}^* = +2.12 \text{ V}$

b.
$$Mg^{2+}(aq) + Ni(s) \rightarrow Mg(s) + Ni^{2+}(aq)$$
 $E_{cell}^* = -2.62 \text{ V}$

c.
$$Mg^{2+}(aq) + Ni^{2+}(aq) \rightarrow Mg(s) + Ni(s)$$
 $E_{cell}^* = +2.62 \text{ V}$

d.
$$Mg(s) + Ni^{2+}(aq) \rightarrow Ni(s) + Mg^{2+}(aq)$$
 $E_{cell}^* = +2.12 \text{ V}$

e.
$$Mg^{2+}(aq) + Ni^{2+}(aq) \rightarrow Ni(s) + Mg(s)$$
 $E_{cell}^* = -2.12 \text{ V}$

19. Calculate E_{cell}^* for the following reaction:

$$F_2(g) + 2 Cl^{-}(aq) \rightarrow 2 F^{-}(aq) + Cl_2(g)$$

given the following standard reduction potentials.

$$F_2(g) + 2 e^- \rightarrow 2 F^-(aq)$$
 $E^0 = +2.87 V$
 $Cl_2(g) + 2 e^- \rightarrow 2 Cl^-(aq)$ $E^0 = +1.36 V$

- a. -4.23 V
- b. -1.51 V
- c. 0.76 V
- d. +1.51 V
- e. +4.23 V
- 20. Calculate E for the following electrochemical cell at 25°C

$$Ag^{+} \mid Ag(aq, 0.150 \text{ M}) \parallel Sn^{2+}(aq, 0.500 \text{ M}), Sn^{4+}(aq, 0.500 \text{ M}) \mid Pt$$

$$Ag^{+}(aq) + e^{-} \rightarrow Ag(s)$$
 $E^{\circ} = +0.80 \text{ V}$
 $Sn^{4+}(aq) + 2 e^{-} \rightarrow Sn^{2+}(aq)$ $E^{\circ} = +0.14 \text{ V}$

- a. -0.915 V
- b. +0.61 V
- c. +0.89 V
- d. +0.915 V
- e. +0.99 V
- _ 21. E_{cell}^* for the following redox reaction is +0.236 V.

$$2 \text{ Fe}^{3+}(aq) + 2 \text{ I}^{-}(aq) \rightarrow 2 \text{ Fe}^{2+}(aq) + \text{I}_{2}(s)$$

What is \triangle °G for this reaction?

- a. -91.0 kJ
- b. -78.1 kJ
- c. -47.2 kJ
- d. -45.5 kJ
- e. -22.8 kJ
- 22. Al³⁺ is reduced to Al(s) at an electrode. If a current of 1.00 ampere is passed for 24 hours, what mass of aluminum is deposited at the electrode? Assume 100 % current efficiency.
 - a. 1.87 g
 - b. 8.05 g
 - c. 24.2 g
 - d. 54.1 g
 - e. 72.5 g
- 23. The solubility of PbCl₂ is 0.016 mol/L. What is the value of $K_{\rm sp}$ for PbCl₂?
 - a. 4.1×10^{-6}
 - b. 1.6×10^{-5}
 - c. 2.6×10^{-4}
 - d. 5.1×10^{-4}
 - e. 4.8×10^{-2}
- The $K_{\rm sp}$ for BaF₂ is 1.7×10^{-6} . What is the concentration of Ba²⁺ in a saturated solution of BaF₂? a. 5.7×10^{-7} M

 - b. $1.7 \times 10^{-6} \text{ M}$
 - c. $1.0 \times 10^{-2} \text{ M}$
 - d. $1.3 \times 10^{-3} \text{ M}$
 - e. $7.5 \times 10^{-3} \text{ M}$
- 25. What is the molar solubility of Ag₂CrO₄ in 0.20 M K₂CrO₄? The value of $K_{\rm sp}$ for Ag₂CrO₄ is 9.0×10^{-12} .

 - c. $3.6 \times 10^{-8} \text{ mol/L}$
 - d. $6.7 \times 10^{-6} \text{ mol/L}$
 - e. $1.3 \times 10^{-4} \text{ mol/L}$
- 26. What is the molar solubility of Fe³⁺ in a solution that is buffered at a pH of 4.00? The $K_{\rm sp}$ for Fe(OH)₃ is 6.3 × 10^{-38} .
 - a. $6.3 \times 10^{-23} \text{ mol/L}$

- b. $2.3 \times 10^{-9} \text{ mol/L}$
- c. $6.3 \times 10^{-8} \text{ mol/L}$
- d. $2.3 \times 10^{-6} \text{ mol/L}$
- e. $1.4 \times 10^{-4} \text{ mol/L}$
- 27. The concentration of Ca²⁺ in a solution is 4.7×10^{-4} M. What concentration of CO₃²⁻ is required to just begin precipitating CaCO₃? The $K_{\rm sp}$ for CaCO₃ is 3.8×10^{-9} .
 - a. $8.1 \times 10^{-6} \text{ M}$
 - b. $5.1 \times 10^{-5} \text{ M}$
 - c. $3.8 \times 10^{-4} \text{ M}$
 - d. $4.7 \times 10^{-4} \text{ M}$
 - e. $1.4 \times 10^{-2} \text{ M}$
 - 28. The following anions can be separated by precipitation as silver salts: Cl⁻, Br⁻, I⁻, CrO₄²⁻. If Ag⁺ is added to a solution containing the four anions, each at a concentration of 0.10 M, in what order would they precipitate?

Compound	K_{sp}
AgCl	1.8×10^{-10}
Ag_2CrO_4	9.0×10^{-12}
AgBr	3.3×10^{-13}
AgI	1.5×10^{-16}

- a. $AgCl \rightarrow Ag_2CrO_4 \rightarrow AgBr \rightarrow AgI$
- b. $AgI \rightarrow AgBr \rightarrow Ag_2CrO_4 \rightarrow AgCl$
- c. $Ag_2CrO_4 \rightarrow AgCl AgBr \rightarrow AgI$
- d. $Ag_2CrO_4 \rightarrow AgI AgBr \rightarrow AgCl$
- e. $AgI \rightarrow AgBr \rightarrow AgCl \rightarrow Ag_2CrO_4$

Essay

29. (6 Pts) Historically, to prevent the oxidation of the iron hulls in ocean vessels, large zinc plates were often affixed to the outside of the hull below the waterline. How does the zinc protect the iron hull?

30. (6 Pts) Explain the function of a salt bridge in a voltaic cell.

Exam3Fall2009thermoelectro Answer Section

MULTIPLE CHOICE

- 1. ANS: C
- 2. ANS: E
- 3. ANS: E
- 4. ANS: A
- 5. ANS: E
- 6. ANS: C
- 7. ANS: B
- 8. ANS: B
- 9. ANS: C
- 10. ANS: B
- 11. ANS: D
- 12. ANS: C
- 13. ANS: A
- 14. ANS: B
- 15. ANS: A
- 16. ANS: E
- 17. ANS: B
- 18. ANS: D
- 19. ANS: D
- 20. ANS: B
- 21. ANS: D
- 22. ANS: B
- 23. ANS: B
- 24. ANS: E
- 25. ANS: D
- 26. ANS: C
- 27. ANS: A
- 28. ANS: E

ESSAY

29. ANS:

The iron, zinc, and seawater form a voltaic cell. The zinc is more easily oxidized than the iron, so it serves as the anode of the cell. Oxidation occurs at the zinc anode, rather than the iron hull.

30. ANS:

The salt bridge physically separates the redox couples in an electrochemical cell while enabling electrical conduction through the cell. In addition, the salt bridge helps maintain electroneutrality as reduction and oxidation occur at the cathode and anode. As oxidation occurs, anions flow from the salt bridge toward the anode and cations flow from the anode to the salt bridge. Likewise, cations flow from the salt bridge toward the cathode and anions from the cathode to the salt bridge.