

3/28/2017

Homework 5

WebAssign

**Homework 5 (Quiz)**

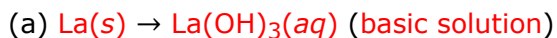
**Current Score** : - / 4

**Due** : Tuesday, April 18 2017 11:59 PM CDT

Bailey Marie Duxworth  
CHEM 1202, section 05, Spring 2017  
Instructor: John Hogan

1. -/0.1 pointsBLB11 20.P.019.

Complete and balance the following half-reactions. In each case indicate whether the half-reaction is an oxidation or reduction. (Use the lowest possible coefficients. Include states-of-matter under SATP conditions in your answer.)




- oxidation
- reduction



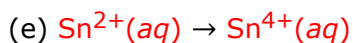

- oxidation
- reduction




- oxidation
- reduction




- oxidation
- reduction

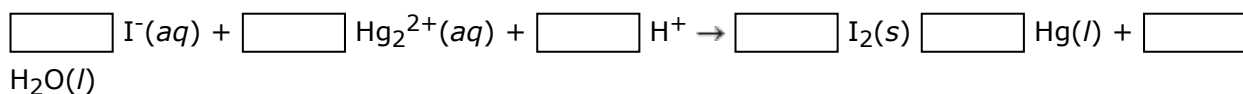



- oxidation
- reduction

## 2. -/0.1 points

For each of the following reactions, balance the chemical equation, calculate the emf, and calculate  $\Delta G^\circ$  at 298 K. (Use the smallest possible coefficients for  $\text{H}_2\text{O}(l)$ ,  $\text{H}^+(aq)$ , and  $\text{HO}^-(aq)$ . These may be zero.)

(a) Aqueous iodide ion is oxidized to  $\text{I}_2(s)$  by  $\text{Hg}_2^{2+}(aq)$ .



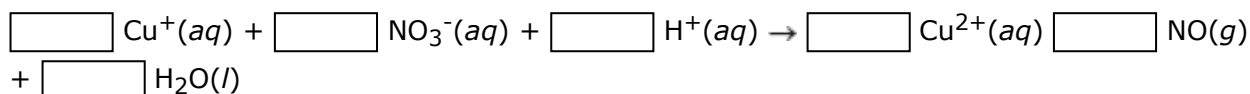
emf

V

$\Delta G^\circ$

kJ

(b) In acidic solution copper(I) ion is oxidized to copper(II) ion by nitrate ion.



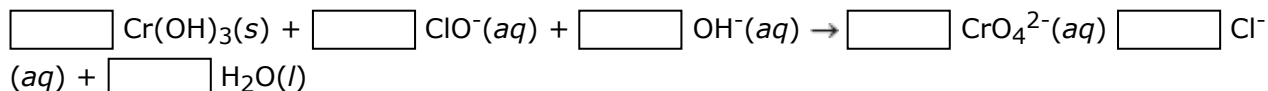
emf

V

$\Delta G^\circ$

kJ

(c) In basic solution  $\text{Cr}(\text{OH})_3(s)$  is oxidized to  $\text{CrO}_4^{2-}(aq)$  by  $\text{ClO}^-(aq)$ .



emf

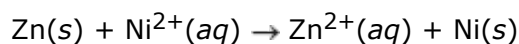
V

$\Delta G^\circ$

kJ

## 3. -/0.1 points

A voltaic cell is constructed that uses the following reaction and operates at 298 K.



(a) What is the emf of this cell under standard conditions?

V

(b) What is the emf of this cell when  $[\text{Ni}^{2+}] = 2.81 \text{ M}$  and  $[\text{Zn}^{2+}] = 0.180 \text{ M}$ ?

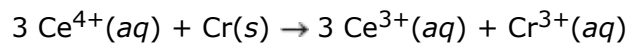
V

(c) What is the emf of the cell when  $[\text{Ni}^{2+}] = 0.288 \text{ M}$  and  $[\text{Zn}^{2+}] = 0.980 \text{ M}$ ?

V

4. -/0.1 points

A voltaic cell utilizes the following reaction and operates at 298 K.



(a) What is the emf of this cell under standard conditions?

V

(b) What is the emf of this cell when  $[\text{Ce}^{4+}] = 1.3 \text{ M}$ ,  $[\text{Ce}^{3+}] = 0.013 \text{ M}$ , and  $[\text{Cr}^{3+}] = 0.014 \text{ M}$ ?

V

(c) What is the emf of the cell when  $[\text{Ce}^{4+}] = 0.54 \text{ M}$ ,  $[\text{Ce}^{3+}] = 0.88 \text{ M}$ , and  $[\text{Cr}^{3+}] = 1.1 \text{ M}$ ?

V

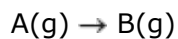
5. -/0.1 points

(a) Select all of the **correct** statements about reaction rates from the choices below.

- The lower the rate of a reaction the longer it takes to reach completion.
- The rate of a fast step has more effect on the overall reaction rate than the rate of a slow step.
- Solid catalysts do not affect reaction rates.
- Solid catalysts increase reaction rates as their surface areas increase.
- The fastest step in a reaction is called the rate-determining step.
- Reaction rates are determined by reactant concentrations, temperatures, and reactant stabilities.
- Reaction rates can show little change as masses of solid reactants increase.

6. -/0.1 points

A flask is charged with 0.150 mol of A and allowed to react to form B according to the following hypothetical gas-phase reaction.



The following data are collected.

times (s)	0	40	80	120	160
moles of A	0.150	0.072	0.049	0.031	0.023

(a) Calculate the number of moles of B at each time in the table.

0 s

mol

40 s

mol

80 s

mol

120 s

mol

160 s

mol

(b) Calculate the average rate of disappearance of A for each 40 s interval, in units of mol/s.

0 - 40 s

mol/s

40 - 80 s

mol/s

80 - 120 s

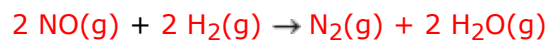
mol/s

120 - 160 mol/s

mol/s

7. -/0.1 points

Consider the following reaction:



(a) The rate law for this reaction is **second** order in **NO(g)** and **first** order in **H<sub>2</sub>(g)**. What is the rate law for this reaction?

- Rate =  $k [\text{NO(g)}] [\text{H}_2\text{(g)}]$
- Rate =  $k [\text{NO(g)}]^2 [\text{H}_2\text{(g)}]$
- Rate =  $k [\text{NO(g)}] [\text{H}_2\text{(g)}]^2$
- Rate =  $k [\text{NO(g)}]^2 [\text{H}_2\text{(g)}]^2$
- Rate =  $k [\text{NO(g)}] [\text{H}_2\text{(g)}]^3$
- Rate =  $k [\text{NO(g)}]^4 [\text{H}_2\text{(g)}]$

(b) If the rate constant for this reaction at a certain temperature is **74300**, what is the reaction rate when **[NO(g)] = 0.0851 M** and **[H<sub>2</sub>(g)] = 0.102 M**?

Rate =  M/s.

(c) What is the reaction rate when the concentration of **NO(g)** is doubled, to **0.170 M** while the concentration of **H<sub>2</sub>(g)** is **0.102 M**?

Rate =  M/s



8. -/0.1 points

The reaction  $2 \text{ClO}_2(\text{aq}) + 2 \text{OH}^-(\text{aq}) \rightarrow \text{ClO}_3^-(\text{aq}) + \text{ClO}_2^-(\text{aq}) + \text{H}_2\text{O}(\text{l})$  was studied at a certain temperature with the following results:

Experiment	$[\text{ClO}_2(\text{aq})]$ (M)	$[\text{OH}^-(\text{aq})]$ (M)	Rate (M/s)
1	0.0494	0.0494	0.0471
2	0.0494	0.0988	0.0943
3	0.0988	0.0494	0.189
4	0.0988	0.0988	0.377

(a) What is the rate law for this reaction?

- Rate =  $k [\text{ClO}_2(\text{aq})] [\text{OH}^-(\text{aq})]$
- Rate =  $k [\text{ClO}_2(\text{aq})]^2 [\text{OH}^-(\text{aq})]$
- Rate =  $k [\text{ClO}_2(\text{aq})] [\text{OH}^-(\text{aq})]^2$
- Rate =  $k [\text{ClO}_2(\text{aq})]^2 [\text{OH}^-(\text{aq})]^2$
- Rate =  $k [\text{ClO}_2(\text{aq})] [\text{OH}^-(\text{aq})]^3$
- Rate =  $k [\text{ClO}_2(\text{aq})]^4 [\text{OH}^-(\text{aq})]$

(b) What is the value of the rate constant?

(c) What is the reaction rate when the concentration of  $\text{ClO}_2(\text{aq})$  is  $0.0929 \text{ M}$  and that of  $\text{OH}^-(\text{aq})$  is  $0.101 \text{ M}$  if the temperature is the same as that used to obtain the data shown above?

M/s

9. -/0.1 points

Select all of the **correct** statements about equilibrium from the choices below.

- At equilibrium the rates of forward and reverse reactions are equal.
- As a reaction proceeds forward toward equilibrium the product concentrations rise.
- As a reaction proceeds forward toward equilibrium the reverse rate constant rises.
- At equilibrium the rate constants of forward and reverse reactions are equal.
- At equilibrium all reactions stop.
- Reactants are transformed into products even at equilibrium.

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10.-/0.1 points

Gaseous  $\text{BrCl}$  is placed in a closed container at  $995\text{ }^\circ\text{C}$ , where it partially decomposes to  $\text{Br}_2$  and  $\text{Cl}_2$ :



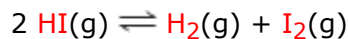
At equilibrium it is found that  $p(\text{BrCl}) = 0.002040$  atm,  $p(\text{Br}_2) = 0.002880$  atm, and  $p(\text{Cl}_2) = 0.005670$  atm. What is the value of  $K_p$  at this temperature?

$K_p =$   .

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11.-/0.1 points

At  $301\text{ }^\circ\text{C}$  the equilibrium constant for the reaction:



is  $K_p = 2.64\text{e-}10$ . If the initial pressure of  $\text{HI}$  is  $0.00247$  atm, what are the equilibrium partial pressures of  $\text{HI}$ ,  $\text{H}_2$ , and  $\text{I}_2$ ?

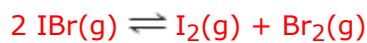
$p(\text{HI}) =$   .

$p(\text{H}_2) =$   .

$p(\text{I}_2) =$   .

12.-/0.1 points

Consider the following equilibrium for which  $\Delta H = -19.51$ :



How will each of the following changes affect an equilibrium mixture of the 3 gases in this reaction?

(a)  $\text{Br}_2(g)$  is added to the system.

---Select---

(b) The reaction mixture is cooled.

---Select---

(c) The volume of the reaction vessel is doubled.

---Select---

(d) A catalyst is added to the reaction mixture.

---Select---

(e) The total pressure of the system is increased by adding a noble gas.

---Select---

(f)  $\text{I}_2(g)$  is removed from the system.

---Select---

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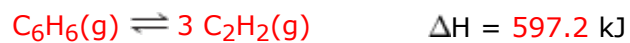
13. -/0.1 points

Choose all of the processes from below which describe changes which are **independent** of the path by which the change occurs.

- the elevation increase experienced by a traveller travelling from Grand Isle, LA to Denver, Colorado
- the kinetic energy acquired by a bullet as it reaches a specific speed
- the work accomplished in burning a gallon of gasoline
- the latitude increase experienced by a traveller travelling from Baton Rouge, LA to Anchorage, Alaska
- the longitude decrease experienced by a traveller travelling from Baton Rouge, LA to London, England
- the work generated by a homogeneous gaseous chemical reaction carried out inside of a bomb calorimeter
- the enthalpy released by the combustion of a gallon of gasoline

14.-/0.1 points

Consider the following reaction which occurs at constant temperature and pressure:



Which has the higher enthalpy,  $\text{C}_6\text{H}_6(\text{g})$  or  $3 \text{C}_2\text{H}_2(\text{g})$ ?

- $\text{C}_6\text{H}_6(\text{g})$
- $3 \text{C}_2\text{H}_2(\text{g})$

Without referring to tables indicate which of each of the pairs of choices below has the higher enthalpy:

(a)

- 1 mol of  $\text{CO}_2(\text{s})$  at 50 K
- 1 mol of  $\text{CO}_2(\text{g})$  at 25 °C

(b)

- 1 mol of  $\text{He}(\text{g})$  at 300 °C
- 1 mol of  $\text{He}(\text{g})$  at 200 °C

(c)

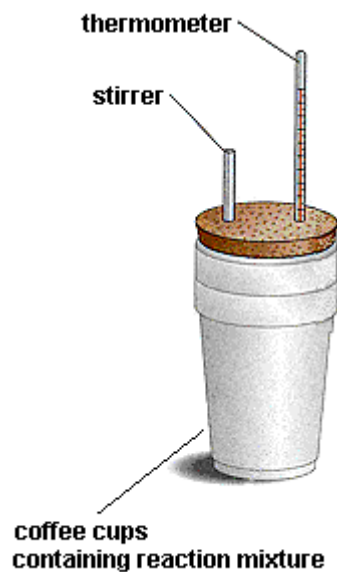
- 1 mol of  $\text{N}_2(\text{g})$  at -100 °C
- 1 mol of  $\text{N}_2(\text{l})$  at -100 °C

(d)

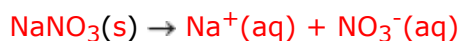
- 1 mol of CO<sub>2</sub>(s) at -78 °C
- 1 mol of CO<sub>2</sub>(g) at -78 °C



15.-/0.1 points



When a 2.74-g sample of solid sodium nitrate dissolves in 41.5 g of water in a coffee-cup calorimeter (see above figure) the temperature falls from 22.00 °C to 18.22 °C. Calculate  $\Delta H$  in kJ/mol  $\text{NaNO}_3$  for the solution process.

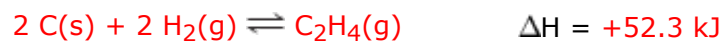
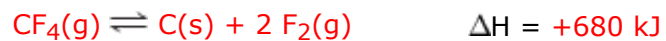
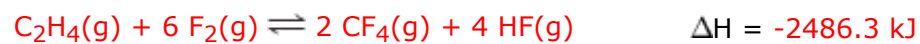


The specific heat of water is 4.18 J/g-K.

$\Delta H_{\text{solution}} =$   kJ/mol  $\text{NaNO}_3$ .

16.-/0.1 points

Calculate the standard enthalpy of formation of gaseous hydrogen fluoride (HF) using the following thermochemical information:



$\Delta\text{H} =$    $\text{ kJ}$

17.-/0.1 points

The normal **condensation** point of **1-propanol gas** is **97 °C**.

(a) Is the **condensation** of **1-propanol gas** an endothermic or an exothermic process?

This process is an  process.

(b) In what temperature range is the **condensation** of **1-propanol gas** a spontaneous process?

This process is spontaneous  **97 °C**.

(c) In what temperature range is this process a nonspontaneous process?

This process is nonspontaneous  **97 °C**.

(d) In what temperature range are the two phases involved in the **condensation** of **1-propanol gas** in equilibrium?

These two phases are in equilibrium  **97 °C**.

18.-/0.1 points

A certain reaction has  $\Delta H^\circ = -80.30$  kJ and  $\Delta S^\circ = 0.00$  J/K.

(a) Is this reaction exothermic, endothermic or isothermic (neither)?

This reaction is  .

(b) Does this reaction lead to a decrease, an increase, or no change in the degree of disorder in the system?

This reaction leads to  in the disorder of the system.

(c) Calculate  $\Delta G^\circ$  for this reaction at 298 K. If this value is less than 1 kJ/mol then enter 0 in the answer box.

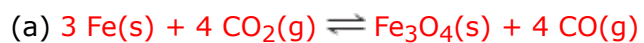
$\Delta G^\circ =$   kJ.

(d) Is this reaction spontaneous, nonspontaneous, or near equilibrium under standard conditions at 298 K?

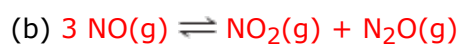
This reaction is  under these conditions.

19.-/0.1 points

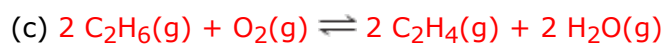
Using values from Appendix C of your textbook, calculate the value of  $K_{eq}$  at 298 K for each of the following reactions:



$K_{eq} =$   .



$K_{eq} =$   .



$K_{eq} =$   .

20.-/0.1 points

For each of the following processes, indicate whether the signs of  $\Delta S$  and  $\Delta H$  are expected to be positive, negative, or about zero.

(a) Ice cubes melt at 5 °C and 1 atm pressure.

For this process  $\Delta S$  should be  and  $\Delta H$  should be .

(b) Liquid water is formed from hydrogen and oxygen gases.

For this process  $\Delta S$  should be  and  $\Delta H$  should be .

(c) A solid sublimes.

For this process  $\Delta S$  should be  and  $\Delta H$  should be .

(d) Pure solid carbon burns in pure oxygen generating carbon dioxide.

For this process  $\Delta S$  should be  and  $\Delta H$  should be .

(e) A nearly ideal gas is allowed to expand and no temperature change occurs.

For this process  $\Delta S$  should be  and  $\Delta H$  should be .

21.-/0.1 points

Complete the following table by calculating the missing entries. In each case indicate whether the solution is acidic or basic.

pH	pOH	[H <sup>+</sup> ]	[OH <sup>-</sup> ]	acidic or basic?
13.32	<input type="text"/>	<input type="text"/> M	<input type="text"/> M	<input type="radio"/> acidic <input type="radio"/> basic
<input type="text"/>	1.57	<input type="text"/> M	<input type="text"/> M	<input type="radio"/> acidic <input type="radio"/> basic
<input type="text"/>	<input type="text"/>	$5.90 \times 10^{-13}$ M	<input type="text"/> M	<input type="radio"/> acidic <input type="radio"/> basic
<input type="text"/>	<input type="text"/>	<input type="text"/> M	$5.70 \times 10^{-6}$ M	<input type="radio"/> acidic <input type="radio"/> basic

22.-/0.1 points

Calculate the pH of each of the following strong acid solutions.

(a) 0.00108 M HI

pH =

(b) 0.266 g of  $\text{HIO}_4$  in 17.0 L of solution

pH =

(c) 13.0 mL of 4.50 M HI diluted to 4.10 L

pH =

(d) a mixture formed by adding 38.0 mL of 0.00351 M HI to 79.0 mL of 0.000840 M  $\text{HIO}_4$

pH =



23. -/0.1 points

Determine the pH of each of the following solutions.

(a) 0.495 M boric acid (weak acid with  $K_a = 5.8e-10$ ).

(b) 0.799 M acetic acid (weak acid with  $K_a = 1.8e-05$ ).

(c) 0.230 M pyridine (weak base with  $K_b = 1.7e-09$ ).

24.-/0.1 points

Calculate the percent ionization of acetic acid ( $\text{HC}_2\text{H}_3\text{O}_2$ ) in solutions of each of the following concentrations ( $K_a = 1.8\text{e-}05$ .)

(a) 0.239 M

%

(b) 0.313 M

%

(c) 0.769 M

%

25.-/0.1 points

(a) Select all of the **correct** statements about the relative acid strengths of pairs of acids from the choices below.

- HCl is a stronger acid than H<sub>2</sub>S because Cl is more electronegative than S.
- HAsO<sub>4</sub><sup>2-</sup> is a stronger acid than H<sub>2</sub>AsO<sub>4</sub><sup>-</sup> because it has more charge (is more unstable).
- H<sub>3</sub>AsO<sub>4</sub> is a stronger acid than H<sub>2</sub>AsO<sub>4</sub><sup>-</sup> because it has more acidic H atoms.
- HBrO<sub>2</sub> is a stronger acid than HBrO<sub>3</sub> because it has fewer oxygens surrounding the central Br atom.
- NH<sub>3</sub> is a stronger acid than H<sub>2</sub>O because N is larger than O.
- HF is a stronger acid than HCl because F is more electronegative than Cl.

26.-/0.1 points

(a) Calculate the percent ionization of  $0.00430 \text{ M}$  hypobromous acid ( $K_a = 2.5e-09$ ).

% ionization =  %

(b) Calculate the percent ionization of  $0.00430 \text{ M}$  hypobromous acid in a solution containing  $0.0260 \text{ M}$  sodium hypobromite.

% ionization =  %

27.-/0.1 points

A buffer solution contains  $0.47$  mol of hypochlorous acid (HClO) and  $0.90$  mol of sodium hypochlorite (NaOCl) in  $3.00$  L.

The  $K_a$  of hypochlorous acid (HClO) is  $K_a = 3e-08$ .

(a) What is the pH of this buffer?

pH =

(b) What is the pH of the buffer after the addition of  $0.39$  mol of NaOH? (assume no volume change)

pH =

(c) What is the pH of the original buffer after the addition of  $0.61$  mol of HI? (assume no volume change)

pH =

28.-/0.1 points

Consider the titration of 80.0 mL of 0.0200 M  $\text{NH}_3$  (a weak base;  $K_b = 1.80 \times 10^{-5}$ ) with 0.100 M  $\text{HIO}_4$ . Calculate the pH after the following volumes of titrant have been added:

(a) 0.0 mL

pH =

(b) 4.0 mL

pH =

(c) 8.0 mL

pH =

(d) 12.0 mL

pH =

(e) 16.0 mL

pH =

(f) 25.6 mL

pH =

29.-/0.1 points

(a) If the molar solubility of  $\text{Cd}_3(\text{PO}_4)_2$  at 25 °C is  $1.19\text{e-}07$  mol/L, what is the  $K_{\text{sp}}$  at this temperature?

$$K_{\text{sp}} = \text{[ ]}$$

(b) It is found that  $0.00100$  g of  $\text{BaCO}_3$  dissolves per 100 mL of aqueous solution at 25 °C. Calculate the solubility-product constant for  $\text{BaCO}_3$ .

$$K_{\text{sp}} = \text{[ ]}$$

(c) The  $K_{\text{sp}}$  of  $\text{Sc}(\text{OH})_3$  at 25 °C is  $2.22\text{e-}31$ . What is the molar solubility of  $\text{Sc}(\text{OH})_3$ ?

$$\text{solubility} = \text{[ ] mol/L}$$

30.-/0.1 points

A solution of  $\text{Na}_2\text{CO}_3$  is added dropwise to a solution that is  $0.0725 \text{ M}$  in  $\text{Nd}^{3+}$  and  $3.05\text{e-}08 \text{ M}$  in  $\text{Hg}_2^{2+}$ .

The  $K_{\text{sp}}$  of  $\text{Nd}_2(\text{CO}_3)_3$  is  $1.08\text{e-}33$ .

The  $K_{\text{sp}}$  of  $\text{Hg}_2\text{CO}_3$  is  $3.6\text{e-}17$ .

(a) What concentration of  $\text{CO}_3^{2-}$  is necessary to begin precipitation? (Neglect volume changes.)

$[\text{CO}_3^{2-}] = \text{[ ]} \text{ M}$ .

(b) Which cation precipitates first?

$\text{Nd}^{3+}$

$\text{Hg}_2^{2+}$

(c) What is the concentration of  $\text{CO}_3^{2-}$  when the second cation begins to precipitate?

$[\text{CO}_3^{2-}] = \text{[ ]} \text{ M}$ .

31.-/0.1 points

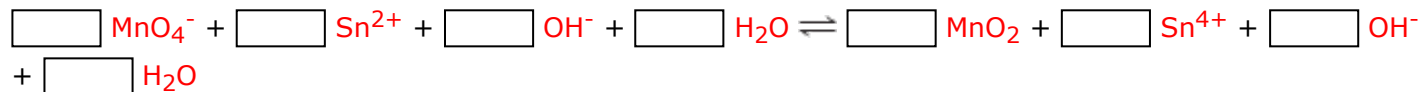
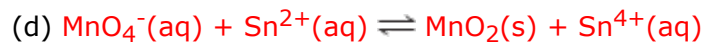
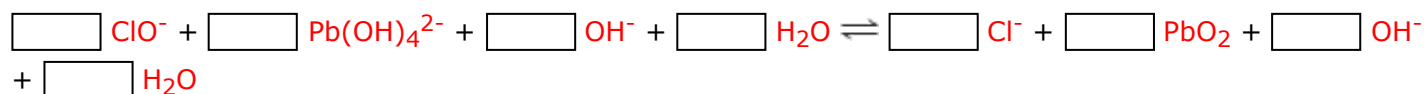
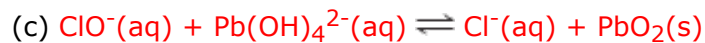
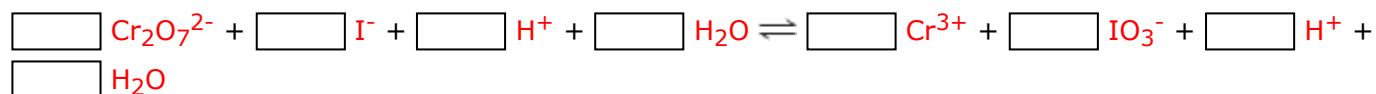
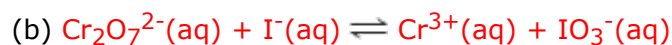
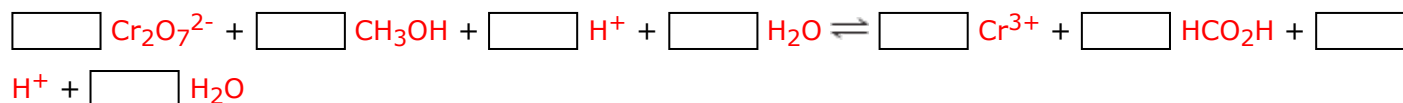
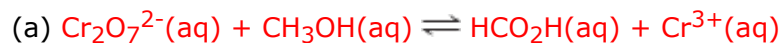
Choose all of the statements from below which are **true** about oxidation and reduction.

- Oxidation and reduction are opposite processes with respect to bookkeeping electrons.
- Oxidation is what happens when an element in a substance gains electrons.
- Electrons appear on the right side of an oxidation half reaction.
- An oxidant is an oxidizing agent, which becomes oxidized during the course of a redox reaction.
- Reduction is what happens when an element in a substance loses electrons.
- Electrons appear on the left side of a reduction half reaction.
- A reductant is an reducing agent, which becomes oxidized during the course of a redox reaction.

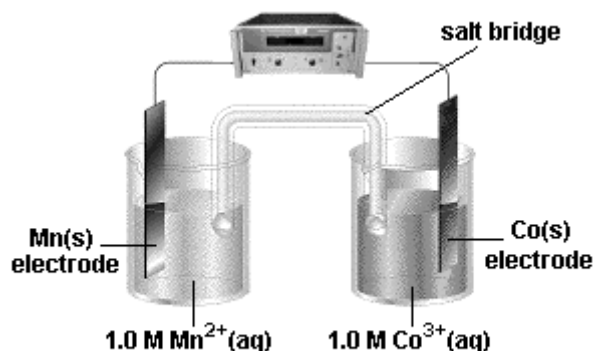


32.-/0.1 points

Balance the following equations. (Use the lowest possible whole-number coefficients. These may be zero.)



33.-/0.1 points



A **voltaic** cell similar to that shown in the figure above is constructed. The electronic device shown at the top of the figure is a **volt meter**. One electrode compartment consists of a **cobalt** strip placed in a **0.5 M**  $\text{Co}_2(\text{SO}_4)_3$  solution, and the other has a **manganese** strip placed in a **1.0 M**  $\text{Mn}(\text{NO}_3)_2$  solution. The overall cell reaction is:



(a) Fill in the information necessary to complete the half reactions that occur in the two electrode compartments. Use the lowest-possible whole-number coefficients. To input an ion surround the ion's element symbol with square brackets and put the ion's charge to the right like this:  $\text{Al}^{3+} = [\text{Al}]3+$  (**not**  $[\text{Al}]+3$ ), and  $\text{Li}^+ = [\text{Li}]+$  (**not**  $[\text{Li}]1+$  or  $[\text{Li}]+1$ ). Do not use brackets for neutral species.

Anode half reaction:  (s)  $\rightleftharpoons$   (aq) +   $\text{e}^-$

Cathode half reaction:  (aq) +   $\text{e}^- \rightleftharpoons$   (s)

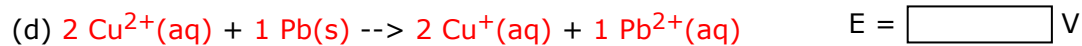
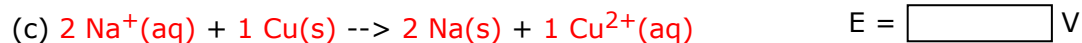
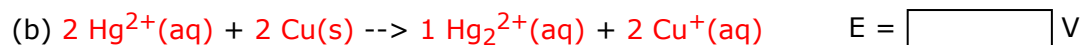
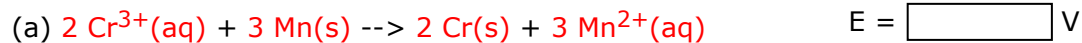
(b) Figure out which electrode is the anode, which is the cathode, the signs on these electrodes, the direction in which the electrons flow through the **volt meter**, and the direction in which cations and anions migrate through the salt bridge and solutions. Use this work to pick all of the statements from below which are **true**.

- Electrons flow from the **manganese** electrode to the **cobalt** electrode through the **volt meter**.

- The **cobalt** electrode is the anode and the **manganese** electrode is the cathode.
- Electrons flow from the **cobalt** electrode to the **manganese** electrode through the **volt meter**.
- The **manganese** electrode is the anode and the **cobalt** electrode is the cathode.
- Anions migrate from the **1.0 M Mn(NO<sub>3</sub>)<sub>2</sub>** solution through the salt bridge to the **0.5 M Co<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>** solution.
- Cations migrate from the **1.0 M Mn(NO<sub>3</sub>)<sub>2</sub>** solution through the salt bridge to the **0.5 M Co<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>** solution.
- Cations migrate from the **0.5 M Co<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>** solution through the salt bridge to the **1.0 M Mn(NO<sub>3</sub>)<sub>2</sub>** solution.
- Anions migrate from the **0.5 M Co<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>** solution through the salt bridge to the **1.0 M Mn(NO<sub>3</sub>)<sub>2</sub>** solution.
- The **manganese** electrode is positive and the **cobalt** electrode is negative.
- The **cobalt** electrode is positive and the **manganese** electrode is negative.

34.-/0.1 points

Using data found in Appendix E of your textbook calculate the nonstandard emf for each of the following reactions if the concentration of each of the ions in these reactions is 0.0009 molar and everything else is standard (use 298 K for the temperature,  $R = 8.314 \text{ J/mol-K}$ , and  $F = 96,485 \text{ C/mol}$ ):



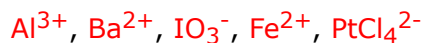
35.-/0.1 points

Choose all of the statements from below which are **true** about strengths of oxidants and reductants.

- The strengths of oxidants and reductants on opposite sides of redox equations correlate oppositely.
- For a strong reductant  $E^{\circ}_{\text{red}}$  should be positive.
- Reducing agents are found on the right-hand side of reduction half reactions.
- For a strong oxidant  $E^{\circ}_{\text{red}}$  should be negative.
- Oxidizing agents are found on the right-hand side of reduction half reactions.

36.-/0.1 points

(a) Assuming standard conditions, arrange the following in order of increasing strength as oxidizing agents in strong acid:



- |                                            |                                            |                                            |                                            |                                            |
|--------------------------------------------|--------------------------------------------|--------------------------------------------|--------------------------------------------|--------------------------------------------|
| <input type="radio"/> $\text{Al}^{3+}$     | <input type="radio"/> $\text{Al}^{3+}$     | <input type="radio"/> $\text{Al}^{3+}$     | <input type="radio"/> $\text{Al}^{3+}$     | <input type="radio"/> $\text{Al}^{3+}$     |
| <input type="radio"/> $\text{Ba}^{2+}$     | <input type="radio"/> $\text{Ba}^{2+}$     | <input type="radio"/> $\text{Ba}^{2+}$     | <input type="radio"/> $\text{Ba}^{2+}$     | <input type="radio"/> $\text{Ba}^{2+}$     |
| <input type="radio"/> $\text{IO}_3^-$      | < <input type="radio"/> $\text{IO}_3^-$    | < <input type="radio"/> $\text{IO}_3^-$    | < <input type="radio"/> $\text{IO}_3^-$    | < <input type="radio"/> $\text{IO}_3^-$    |
| <input type="radio"/> $\text{Fe}^{2+}$     | <input type="radio"/> $\text{Fe}^{2+}$     | <input type="radio"/> $\text{Fe}^{2+}$     | <input type="radio"/> $\text{Fe}^{2+}$     | <input type="radio"/> $\text{Fe}^{2+}$     |
| <input type="radio"/> $\text{PtCl}_4^{2-}$ | <input type="radio"/> $\text{PtCl}_4^{2-}$ | <input type="radio"/> $\text{PtCl}_4^{2-}$ | <input type="radio"/> $\text{PtCl}_4^{2-}$ | <input type="radio"/> $\text{PtCl}_4^{2-}$ |

(b) Assuming standard conditions, arrange the following in order of increasing strength as reducing agents in strong base:



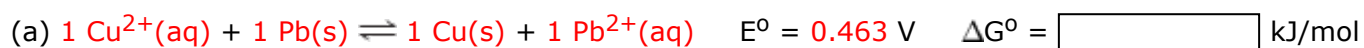
- |                                                |                                                |                                                |                                                |                                                |
|------------------------------------------------|------------------------------------------------|------------------------------------------------|------------------------------------------------|------------------------------------------------|
| <input type="radio"/> NO                       | <input type="radio"/> NO                       | <input type="radio"/> NO                       | <input type="radio"/> NO                       | <input type="radio"/> NO                       |
| <input type="radio"/> $\text{Co}^{2+}$         | <input type="radio"/> $\text{Co}^{2+}$         | <input type="radio"/> $\text{Co}^{2+}$         | <input type="radio"/> $\text{Co}^{2+}$         | <input type="radio"/> $\text{Co}^{2+}$         |
| <input type="radio"/> Cr                       | < <input type="radio"/> Cr                     | < <input type="radio"/> Cr                     | < <input type="radio"/> Cr                     | < <input type="radio"/> Cr                     |
| <input type="radio"/> $\text{H}_3\text{AsO}_3$ | <input type="radio"/> $\text{H}_3\text{AsO}_3$ | <input type="radio"/> $\text{H}_3\text{AsO}_3$ | <input type="radio"/> $\text{H}_3\text{AsO}_3$ | <input type="radio"/> $\text{H}_3\text{AsO}_3$ |
| <input type="radio"/> $\text{Br}^-$            | <input type="radio"/> $\text{Br}^-$            | <input type="radio"/> $\text{Br}^-$            | <input type="radio"/> $\text{Br}^-$            | <input type="radio"/> $\text{Br}^-$            |

37.-/0.1 points

(a) Choose all of the statements from below which **correctly** describe issues pertaining to the relationship between the emf of a reaction and its Gibbs free energy change.

- The Gibbs free energy change and emf values of a reaction have opposite algebraic signs (+ vs. -).
- Faraday's constant is used to interconvert coulombs of electrons and moles of electrons.
- Electron stoichiometry need not be used to interconvert a Gibbs free energy change and a reaction emf.
- A Gibbs free energy change involves moles of substances whereas an emf involves coulombs of electrons.
- Electron stoichiometry must be used to interconvert a Gibbs free energy change and a reaction emf.

Calculate the standard Gibbs free energy changes at 25 °C for each of the reactions shown below using the  $E^\circ$  values given. Select whether each of these reactions is nonspontaneous, at equilibrium, or spontaneous under standard conditions.



- nonspontaneous
- at equilibrium
- spontaneous

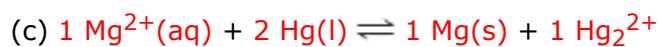


(aq)

V

kJ/mol

- nonspontaneous
- at equilibrium
- spontaneous



(aq)

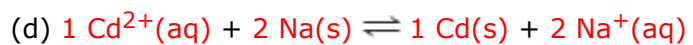
$$E^\circ = -3.159$$

V

$$\Delta G^\circ =$$

kJ/mol

- nonspontaneous
- at equilibrium
- spontaneous



$$E^\circ = 2.307 \text{ V}$$

$$\Delta G^\circ =$$

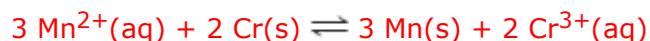
kJ/mol

- nonspontaneous
- at equilibrium
- spontaneous



38.-/0.1 points

A **electrolytic** cell operating under **standard conditions** (1.0 M ion concentrations) utilizes the following reaction:



What is the effect on the cell emf of each of the following changes?

(a) Water is added to the **anode** compartment, diluting the solution.

- The positive cell emf rises, becoming more positive.
- The positive cell emf drops closer to zero.
- The negative cell emf rises closer to zero.
- The negative cell emf drops, becoming more negative.
- No change in cell emf occurs.

(b) The size of the **manganese** electrode is increased.

- The positive cell emf rises, becoming more positive.
- The positive cell emf drops closer to zero.
- The negative cell emf rises closer to zero.
- The negative cell emf drops, becoming more negative.
- No change in cell emf occurs.

(c) A solution of 1.0 M **MnCl<sub>2</sub>** is added to the **cathode** compartment.

- The positive cell emf rises, becoming more positive.
- The positive cell emf drops closer to zero.
- The negative cell emf rises closer to zero.
- The negative cell emf drops, becoming more negative.

No change in cell emf occurs.

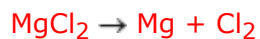
(d) Some  $\text{Na}_2\text{S}$  is added to the  $\text{Cr}^{3+}$  compartment, precipitating some  $\text{Cr}^{3+}$  as chromium sulfide.

- The positive cell emf rises, becoming more positive.
- The positive cell emf drops closer to zero.
- The negative cell emf rises closer to zero.
- The negative cell emf drops, becoming more negative.
- No change in cell emf occurs.

39.-/0.1 points

Metallic magnesium can be made by the electrolysis of molten  $\text{MgCl}_2$ .

(a) What mass of  $\text{Mg}$  is formed by passing a current of 8.39 A through molten  $\text{MgCl}_2$  for 2.80 days? The unbalanced chemical reaction representing this electrolysis is shown below.



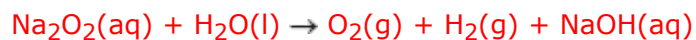
g of  $\text{Mg}$  is formed by this electrolysis.

(b) How many minutes are needed to plate out 6.00 g of  $\text{Mg}$  from molten  $\text{MgCl}_2$  using 5.14 A current?

minutes are needed.

40.-/0.1 points

(a) In the electrolysis of aqueous  $\text{Na}_2\text{O}_2$ , how many liters of  $\text{H}_2(\text{g})$  (at STP) are generated by a current of 64.5 A for a period of 58.3 min? The unbalanced chemical reaction representing this electrolysis is shown below.



liters of  $\text{H}_2(\text{g})$  is generated by this electrolysis.

(b) How many moles of  $\text{NaOH}(\text{aq})$  are formed in the solution in this process?

moles of  $\text{NaOH}(\text{aq})$  are formed.