AP · ⊖ CollegeBoard

AP Chemistry
Free-Response Questions

PERIODIC TABLE OF THE ELEMENTS

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18
1 **H** 1.008																	2 **He** 4.00
3 **Li** 6.94	4 **Be** 9.01											5 **B** 10.81	6 **C** 12.01	7 **N** 14.01	8 **O** 16.00	9 **F** 19.00	10 **Ne** 20.18
11 **Na** 22.99	12 **Mg** 24.30											13 **Al** 26.98	14 **Si** 28.09	15 **P** 30.97	16 **S** 32.06	17 **Cl** 35.45	18 **Ar** 39.95
19 **K** 39.10	20 **Ca** 40.08	21 **Sc** 44.96	22 **Ti** 47.87	23 **V** 50.94	24 **Cr** 52.00	25 **Mn** 54.94	26 **Fe** 55.85	27 **Co** 58.93	28 **Ni** 58.69	29 **Cu** 63.55	30 **Zn** 65.38	31 **Ga** 69.72	32 **Ge** 72.63	33 **As** 74.92	34 **Se** 78.97	35 **Br** 79.90	36 **Kr** 83.80
37 **Rb** 85.47	38 **Sr** 87.62	39 **Y** 88.91	40 **Zr** 91.22	41 **Nb** 92.91	42 **Mo** 95.95	43 **Tc** (97)	44 **Ru** 101.1	45 **Rh** 102.91	46 **Pd** 106.42	47 **Ag** 107.87	48 **Cd** 112.41	49 **In** 114.82	50 **Sn** 118.71	51 **Sb** 121.76	52 **Te** 127.60	53 **I** 126.90	54 **Xe** 131.29
55 **Cs** 132.91	56 **Ba** 137.33	57 ***La** 138.91	72 **Hf** 178.49	73 **Ta** 180.95	74 **W** 183.84	75 **Re** 186.21	76 **Os** 190.2	77 **Ir** 192.2	78 **Pt** 195.08	79 **Au** 196.97	80 **Hg** 200.59	81 **Tl** 204.38	82 **Pb** 207.2	83 **Bi** 208.98	84 **Po** (209)	85 **At** (210)	86 **Rn** (222)
87 **Fr** (223)	88 **Ra** (226)	89 **†Ac** (227)	104 **Rf** (267)	105 **Db** (270)	106 **Sg** (271)	107 **Bh** (270)	108 **Hs** (277)	109 **Mt** (276)	110 **Ds** (281)	111 **Rg** (282)	112 **Cn** (285)	113 **Uut** (285)	114 **Fl** (289)	115 **Uup** (288)	116 **Lv** (293)	117 **Uus** (294)	118 **Uuo** (294)

*Lanthanoid Series

58 **Ce** 140.12	59 **Pr** 140.91	60 **Nd** 144.24	61 **Pm** (145)	62 **Sm** 150.4	63 **Eu** 151.97	64 **Gd** 157.25	65 **Tb** 158.93	66 **Dy** 162.50	67 **Ho** 164.93	68 **Er** 167.26	69 **Tm** 168.93	70 **Yb** 173.05	71 **Lu** 174.97

† Actinoid Series

90 **Th** 232.04	91 **Pa** 231.04	92 **U** 238.03	93 **Np** (237)	94 **Pu** (244)	95 **Am** (243)	96 **Cm** (247)	97 **Bk** (247)	98 **Cf** (251)	99 **Es** (252)	100 **Fm** (257)	101 **Md** (258)	102 **No** (259)	103 **Lr** (262)

AP® CHEMISTRY EQUATIONS AND CONSTANTS

Throughout the exam the following symbols have the definitions specified unless otherwise noted.

L, mL	=	liter(s), milliliter(s)	mm Hg	=	millimeters of mercury
g	=	gram(s)	J, kJ	=	joule(s), kilojoule(s)
nm	=	nanometer(s)	V	=	volt(s)
atm	=	atmosphere(s)	mol	=	mole(s)

ATOMIC STRUCTURE

$$E = h\nu$$
$$c = \lambda\nu$$

E = energy
ν = frequency
λ = wavelength

Planck's constant, $h = 6.626 \times 10^{-34}$ J s
Speed of light, $c = 2.998 \times 10^8$ m s^{-1}
Avogadro's number $= 6.022 \times 10^{23}$ mol^{-1}
Electron charge, $e = -1.602 \times 10^{-19}$ coulomb

EQUILIBRIUM

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}, \text{ where } a\,A + b\,B \rightleftarrows c\,C + d\,D$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][HB^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} \text{ at } 25°C$$
$$= K_a \times K_b$$

$$pH = -\log[H^+], \quad pOH = -\log[OH^-]$$

$$14 = pH + pOH$$

$$pH = pK_a + \log\frac{[A^-]}{[HA]}$$

$$pK_a = -\log K_a, \quad pK_b = -\log K_b$$

Equilibrium Constants

K_c (molar concentrations)
K_p (gas pressures)
K_a (weak acid)
K_b (weak base)
K_w (water)

KINETICS

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\frac{1}{[A]_t} - \frac{1}{[A]_0} = kt$$

$$t_{1/2} = \frac{0.693}{k}$$

k = rate constant
t = time
$t_{1/2}$ = half-life

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$P_A = P_{total} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{total} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = {}^\circ C + 273$$

$$D = \frac{m}{V}$$

$$KE \text{ per molecule} = \frac{1}{2}mv^2$$

Molarity, M = moles of solute per liter of solution

$$A = abc$$

P = pressure
V = volume
T = temperature
n = number of moles
m = mass
M = molar mass
D = density
KE = kinetic energy
v = velocity
A = absorbance
a = molar absorptivity
b = path length
c = concentration

Gas constant, R = 8.314 J mol^{-1} K^{-1}

= 0.08206 L atm mol^{-1} K^{-1}

= 62.36 L torr mol^{-1} K^{-1}

1 atm = 760 mm Hg = 760 torr

STP = 273.15 K and 1.0 atm

Ideal gas at STP = 22.4 L mol^{-1}

THERMODYNAMICS / ELECTROCHEMISTRY

$$q = mc\Delta T$$

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

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

$$= -RT \ln K$$

$$= -nFE^\circ$$

$$I = \frac{q}{t}$$

q = heat
m = mass
c = specific heat capacity
T = temperature
S° = standard entropy
H° = standard enthalpy
G° = standard Gibbs free energy
n = number of moles
E° = standard reduction potential
I = current (amperes)
q = charge (coulombs)
t = time (seconds)

Faraday's constant, F = 96,485 coulombs per mole of electrons

$$1 \text{ volt} = \frac{1 \text{ joule}}{1 \text{ coulomb}}$$

CHEMISTRY
Section II
Time—1 hour and 45 minutes
7 Questions

YOU MAY USE YOUR CALCULATOR FOR THIS SECTION.

Directions: Questions 1–3 are long free-response questions that require about 23 minutes each to answer and are worth 10 points each. Questions 4–7 are short free-response questions that require about 9 minutes each to answer and are worth 4 points each.

Write your response in the space provided following each question. Examples and equations may be included in your responses where appropriate. For calculations, clearly show the method used and the steps involved in arriving at your answers. You must show your work to receive credit for your answer. Pay attention to significant figures.

$$Na_2S_2O_3(aq) + 4\, NaOCl(aq) + 2\, NaOH(aq) \rightarrow 2\, Na_2SO_4(aq) + 4\, NaCl(aq) + H_2O(l)$$
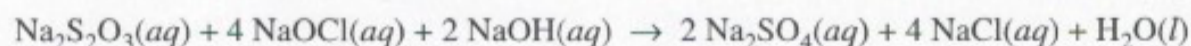
1. A student performs an experiment to determine the value of the enthalpy change, ΔH°_{rxn}, for the oxidation-reduction reaction represented by the balanced equation above.

   (a) Determine the oxidation number of Cl in NaOCl.

   (b) Calculate the number of grams of $Na_2S_2O_3$ needed to prepare 100.00 mL of 0.500 M $Na_2S_2O_3(aq)$.

   In the experiment, the student uses the solutions shown in the table below.

Solution	Concentration (M)	Volume (mL)
$Na_2S_2O_3(aq)$	0.500	5.00
$NaOCl(aq)$	0.500	5.00
$NaOH(aq)$	0.500	5.00

   (c) Using the balanced equation for the oxidation-reduction reaction and the information in the table above, determine which reactant is the limiting reactant. Justify your answer.

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The solutions, all originally at 20.0°C, are combined in an insulated calorimeter. The temperature of the reaction mixture is monitored, as shown in the graph below.
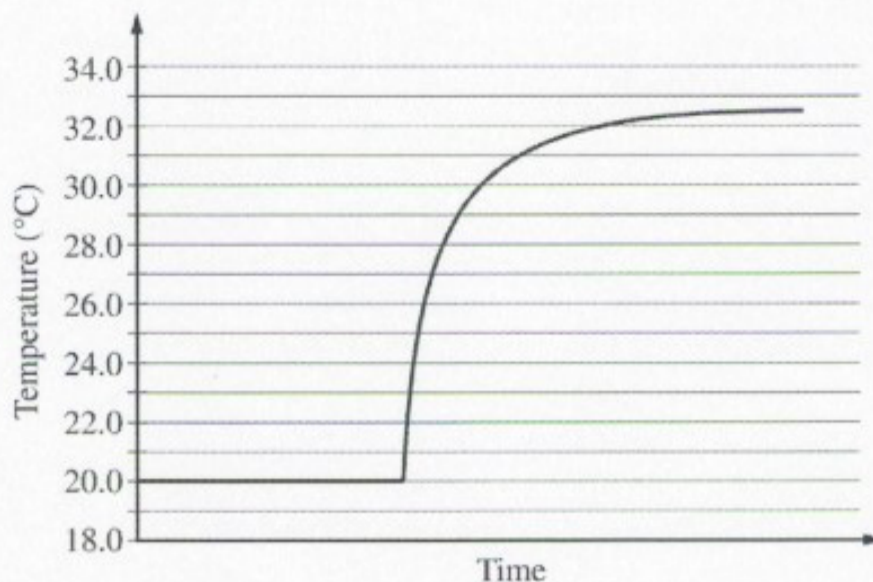
(d) According to the graph, what is the temperature change of the reaction mixture?

(e) The mass of the reaction mixture inside the calorimeter is 15.21 g.

   (i) Calculate the magnitude of the heat energy, in joules, that is released during the reaction. Assume that the specific heat of the reaction mixture is 3.94 J/(g·°C) and that the heat absorbed by the calorimeter is negligible.

   (ii) Using the balanced equation for the oxidation-reduction reaction and your answer to part (c), calculate the value of the enthalpy change of the reaction, ΔH°_{rxn}, in kJ/mol$_{rxn}$. Include the appropriate algebraic sign with your answer.

The student repeats the experiment, but this time doubling the volume of each of the reactants, as shown in the table below.

Solution	Concentration (M)	Volume (mL)
$Na_2S_2O_3(aq)$	0.500	10.0
$NaOCl(aq)$	0.500	10.0
$NaOH(aq)$	0.500	10.0

(f) The magnitude of the enthalpy change, ΔH°_{rxn}, in kJ/mol$_{rxn}$, calculated from the results of the second experiment is the same as the result calculated in part (e)(ii). Explain this result.

(g) Write the balanced net ionic equation for the given reaction.

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$$2\,NO(g) + O_2(g) \rightarrow 2\,NO_2(g)$$
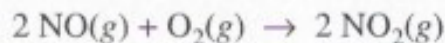
2. A student investigates the reactions of nitrogen oxides. One of the reactions in the investigation requires an equimolar mixture of $NO(g)$ and $NO_2(g)$, which the student produces by using the reaction represented above.

(a) The particle-level representation of the equimolar mixture of $NO(g)$ and $NO_2(g)$ in the flask at the completion of the reaction between $NO(g)$ and $O_2(g)$ is shown below in the box on the right. In the box below on the left, draw the particle-level representation of the <u>reactant</u> mixture of $NO(g)$ and $O_2(g)$ that would yield the product mixture shown in the box on the right. In your drawing, represent oxygen atoms and nitrogen atoms as indicated below.

Oxygen atom = ◯   Nitrogen atom = ⬤

Reactant Mixture   Product Mixture
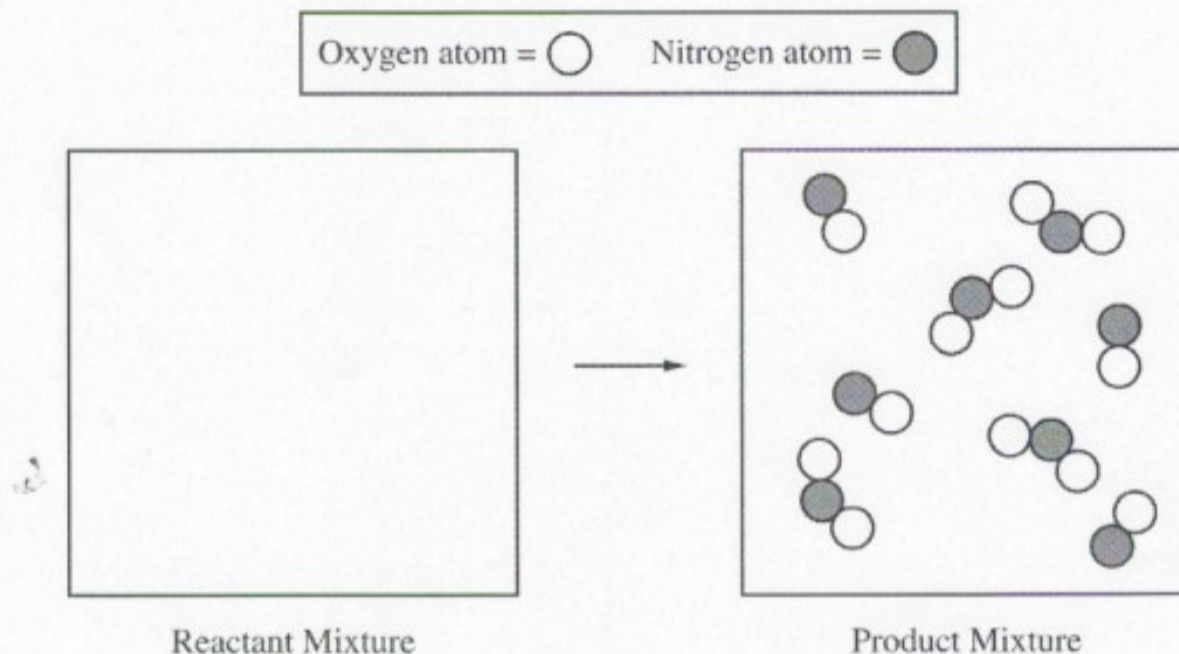
The student reads in a reference text that $NO(g)$ and $NO_2(g)$ will react as represented by the equation below. Thermodynamic data for the reaction are given in the table below the equation.

$$NO(g) + NO_2(g) \rightleftarrows N_2O_3(g)$$
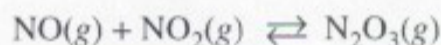
ΔH°_{298}	ΔS°_{298}	ΔG°_{298}
-40.4 kJ/mol$_{rxn}$	-138.5 J/(K·mol$_{rxn}$)	0.87 kJ/mol$_{rxn}$

(b) The student begins with an equimolar mixture of $NO(g)$ and $NO_2(g)$ in a rigid reaction vessel and the mixture reaches equilibrium at 298 K.

   (i) Calculate the value of the equilibrium constant, K, for the reaction at 298 K.

   (ii) If both P_{NO} and P_{NO_2} in the vessel are initially 1.0 atm, will $P_{N_2O_3}$ at equilibrium be equal to 1.0 atm? Justify your answer.

(c) The student hypothesizes that increasing the temperature will increase the amount of $N_2O_3(g)$ in the equilibrium mixture. Indicate whether you agree or disagree with the hypothesis. Justify your answer.

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$N_2O_3(g)$ reacts with water to form nitrous acid, $HNO_2(aq)$, a compound involved in the production of acid rain. The reaction is represented below.

$$N_2O_3(g) + H_2O(l) \rightarrow 2\, HNO_2(aq)$$
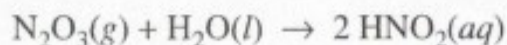
(d) The skeletal structure of the HNO_2 molecule is shown in the box below.

(i) Complete the Lewis electron-dot diagram of the HNO_2 molecule in the box below, including any lone pairs of electrons.

H   O   N   O

(ii) Based on your completed diagram above, identify the hybridization of the nitrogen atom in the HNO_2 molecule.

To produce an aqueous solution of HNO_2, the student bubbles $N_2O_3(g)$ into distilled water. Assume that the reaction goes to completion and that HNO_2 is the only species produced. To determine the concentration of $HNO_2(aq)$ in the resulting solution, the student titrates a 100. mL sample of the solution with 0.100 M KOH(aq). The neutralization reaction is represented below.

$$HNO_2(aq) + OH^-(aq) \rightarrow NO_2^-(aq) + H_2O(l)$$
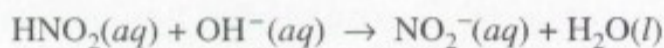
The following titration curve shows the change in pH of the solution during the titration.

Volume of 0.100 M KOH(aq) Added (mL)
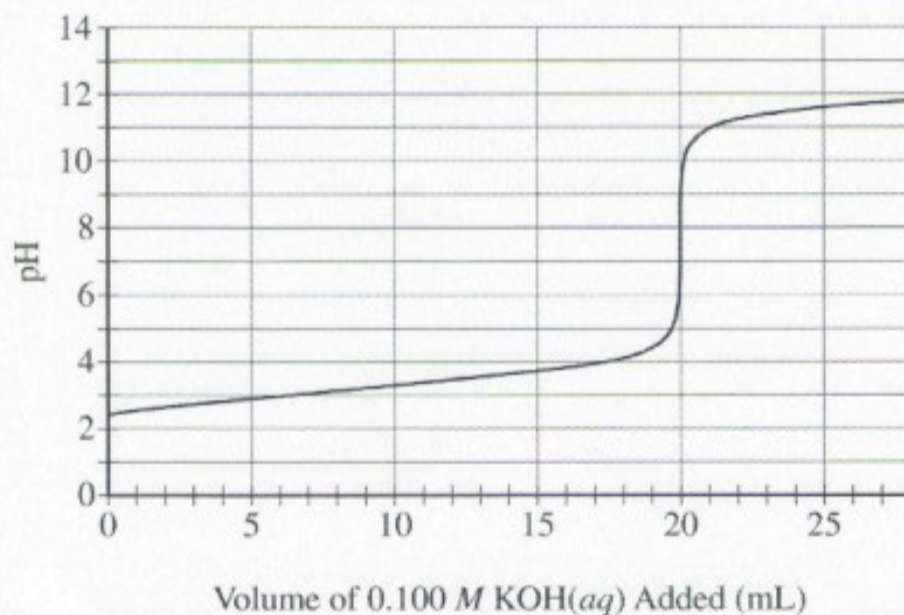
(e) Use the titration curve and the information above to

(i) determine the initial concentration of the $HNO_2(aq)$ solution

(ii) estimate the value of pK_a for $HNO_2(aq)$

(f) During the titration, after a volume of 15 mL of 0.100 M KOH(aq) has been added, which species, $HNO_2(aq)$ or $NO_2^-(aq)$, is present at a higher concentration in the solution? Justify your answer.

3. Answer the following questions relating to Fe and its ions, Fe^{2+} and Fe^{3+}.

   (a) Write the ground-state electron configuration of the Fe^{2+} ion.

Ion	Ionic Radius (pm)
Fe^{2+}	92
Fe^{3+}	79

   (b) The radii of the ions are given in the table above. Using principles of atomic structure, explain why the radius of the Fe^{2+} ion is larger than the radius of the Fe^{3+} ion.

   (c) Fe^{3+} ions interact more strongly with water molecules in aqueous solution than Fe^{2+} ions do. Give one reason for this stronger interaction, and justify your answer using Coulomb's law.

A student obtains a solution that contains an unknown concentration of $Fe^{2+}(aq)$. To determine the concentration of $Fe^{2+}(aq)$ in the solution, the student titrates a sample of the solution with $MnO_4^-(aq)$, which converts $Fe^{2+}(aq)$ to $Fe^{3+}(aq)$, as represented by the following equation.

$$5\ Fe^{2+}(aq) + MnO_4^-(aq) + 8\ H^+(aq) \rightarrow 5\ Fe^{3+}(aq) + Mn^{2+}(aq) + 4\ H_2O(l)$$
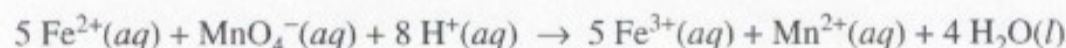
   (d) Write the balanced equation for the half-reaction for the oxidation of $Fe^{2+}(aq)$ to $Fe^{3+}(aq)$.

   (e) The student titrates a 10.0 mL sample of the $Fe^{2+}(aq)$ solution. Calculate the value of $[Fe^{2+}]$ in the solution if it takes 17.48 mL of added 0.0350 M $KMnO_4(aq)$ to reach the equivalence point of the titration.

To deliver the 10.0 mL sample of the $Fe^{2+}(aq)$ solution in part (e), the student has the choice of using one of the pieces of glassware listed below.

   - 25 mL buret
   - 25 mL graduated cylinder
   - 25 mL beaker
   - 25 mL volumetric flask

   (f) Explain why the 25 mL volumetric flask would be a poor choice to use for delivering the required volume of the $Fe^{2+}(aq)$ solution.

In a separate experiment, the student is given a sample of powdered Fe(s) that contains an inert impurity. The student uses a procedure to oxidize the Fe(s) in the sample to $Fe_2O_3(s)$. The student collects the following data during the experiment.

Mass of Fe(s) with inert impurity	6.724 g
Mass of $Fe_2O_3(s)$ produced	7.531 g

   (g) Calculate the number of moles of Fe in the $Fe_2O_3(s)$ produced.

   (h) Calculate the percent by mass of Fe in the original sample of powdered Fe(s) with the inert impurity.

   (i) If the oxidation of the Fe(s) in the original sample was incomplete so that some of the 7.531 g of product was FeO(s) instead of $Fe_2O_3(s)$, would the calculated mass percent of Fe(s) in the original sample be higher, lower, or the same as the actual mass percent of Fe(s)? Justify your answer.

GO ON TO THE NEXT PAGE.

| Sulfur atom = ◯ | Carbon atom = ⬤ | Oxygen atom = ◯ |

Compound	Molecular Structure	Boiling Point at 1 atm (K)
CS_2	◯–⬤–◯	319
COS	◯–⬤–◯	223

4. The table above gives the molecular structures and boiling points for the compounds CS_2 and COS.

(a) In terms of the types and relative strengths of all the intermolecular forces in each compound, explain why the boiling point of $CS_2(l)$ is higher than that of COS(l).

(b) A 10.0 g sample of $CS_2(l)$ is put in an evacuated 5.0 L rigid container. The container is sealed and heated to 325 K, at which temperature all of the $CS_2(l)$ has vaporized. What is the pressure in the container once all of the $CS_2(l)$ has vaporized?

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$$HF(aq) + H_2O(l) \;\rightleftarrows\; F^-(aq) + H_3O^+(aq)$$

5. The ionization of HF(aq) in water is represented by the equation above. In a 0.0350 M HF(aq) solution, the percent ionization of HF is 13.0 percent.

(a) Two particulate representations of the ionization of HF molecules in the 0.0350 M HF(aq) solution are shown below in Figure 1 and Figure 2. Water molecules are not shown. Explain why the representation of the ionization of HF molecules in water in Figure 1 is more accurate than the representation in Figure 2. (The key below identifies the particles in the representations.)
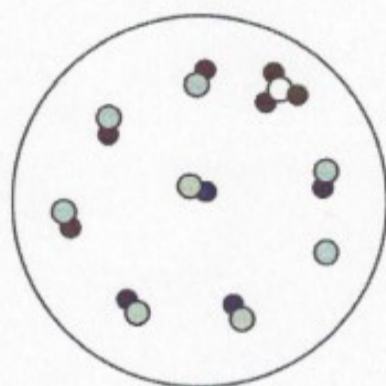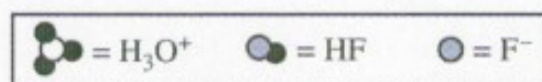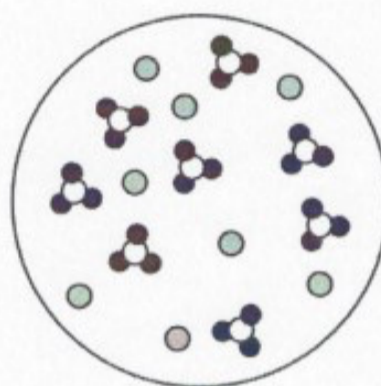
Figure 1   Figure 2

(b) Use the percent ionization data above to calculate the value of K_a for HF.

(c) If 50.0 mL of distilled water is added to 50.0 mL of 0.035 M HF(aq), will the percent ionization of HF(aq) in the solution increase, decrease, or remain the same? Justify your answer with an explanation or calculation.

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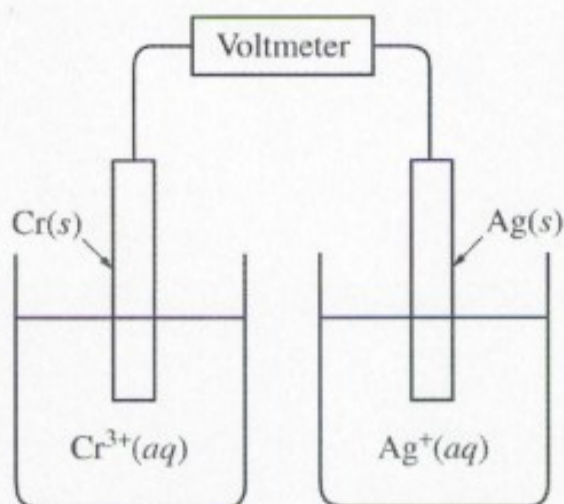
6. A student sets up a galvanic cell at 298 K that has an electrode of Ag(s) immersed in a 1.0 M solution of $Ag^+(aq)$ and an electrode of Cr(s) immersed in a 1.0 M solution of $Cr^{3+}(aq)$, as shown in the diagram above.

(a) The student measures the voltage of the cell shown above and discovers that it is zero. Identify the missing component of the cell, and explain its importance for obtaining a nonzero voltage.

Half-Reaction	$E°$ (V)
$Ag^+(aq) + e^- \rightarrow Ag(s)$	+ 0.80
$Cr^{3+}(aq) + 3\,e^- \rightarrow Cr(s)$	?

(b) The student adds the missing component to the cell and measures $E°_{cell}$ to be +1.54 V. As the cell operates, Ag^+ ions are reduced. Use this information and the information in the table above to do the following.

  (i) Calculate the value of $E°$ for the half-reaction $Cr^{3+}(aq) + 3\,e^- \rightarrow Cr(s)$.

  (ii) Write the balanced net-ionic equation for the overall reaction that occurs as the cell operates.

 (iii) Calculate the value of $\Delta G°$ for the overall cell reaction in J/mol_{rxn}.

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Relative Number of Electrons

Binding Energy (MJ/mol)
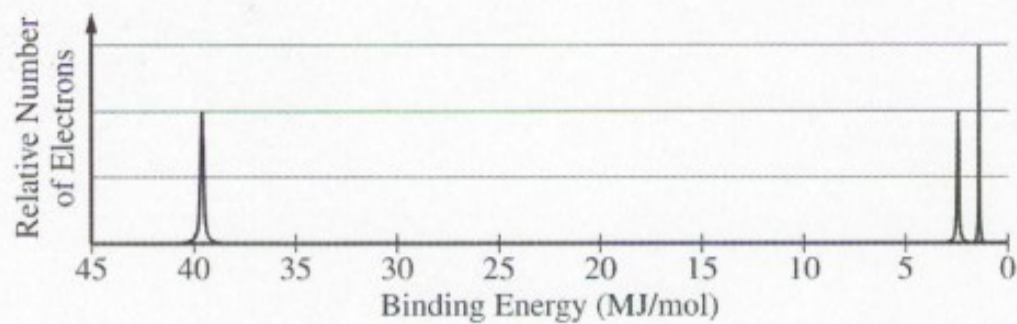
7. The complete photoelectron spectrum of an element is represented above.

  (a) Identify the element.

A radioactive isotope of the element decays with a half-life of 10. minutes.

  (b) Calculate the value of the rate constant, k, for the radioactive decay. Include units with your answer.

  (c) If 64 atoms of the radioactive isotope are originally present in a sample, what is the expected amount of time that will pass until only one atom of the isotope remains? Show how you arrived at your answer.

STOP

END OF EXAM

1)

a. $\overset{+1}{Na} \overset{-2}{O} \overset{+1}{Cl}$ +1
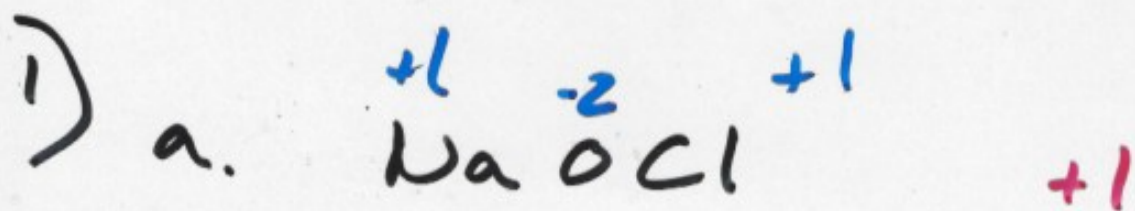
b. $0.1000 L \times \dfrac{0.500 mol \ Na_2 S_2 O_3}{L}$

$\times \dfrac{158.1 g \ Na_2 S_2 O_3}{1 mol} = 7.90 g \ Na_2 S_2 O_3$

c. $Na_2 S_2 O_3 + 4 NaOCl + 2 NaOH$

$\longrightarrow 2 Na_2 SO_4 + 4 NaCl + H_2O$
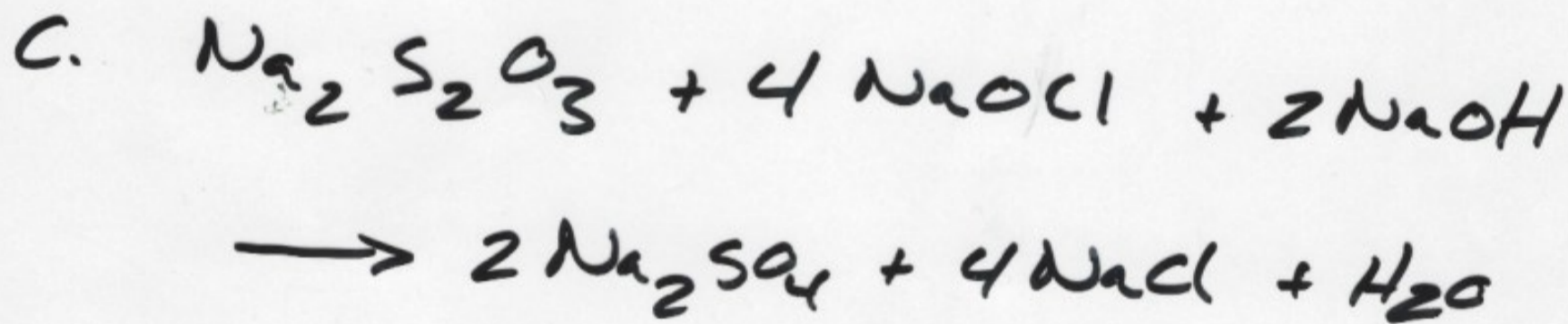
Given equal mole values for the reactants, NaOCl will be limiting. According to the balanced equation for every 1 mol of $Na_2 S_2 O_3$ and for every 2 moles of NaOH 4 moles of NaOCl are required.

1] d. $\Delta T = T_f - T_i$

              $= 32.5°C - 20.0C°$

              $= 12.5°C$

e.

i

   $q = mc\Delta T$

      $= (15.21g)\left(3.94\dfrac{J}{g°C}\right)(12.5°C)$

      $= 749 J$

e ii

   $5.00 mL\ NaOH \times \dfrac{1L}{1000mL}$

$\times \dfrac{0.500 mol}{1L} = 0.0025\ mol\ NaOH$

$\dfrac{749\ J}{0.00250 mol} \times \dfrac{1 KJ}{1000 J} \times \dfrac{4\ mol\ NaOCl}{1 mol\ rxn}$

      $-1.20 \times 10^3 KJ/mol$

1) f. Twice the heat is release for twice the amount of limiting reactant used

$$\frac{2(-1.20 \times 10^{-3} J)}{2 \, mol \, NaOH}$$

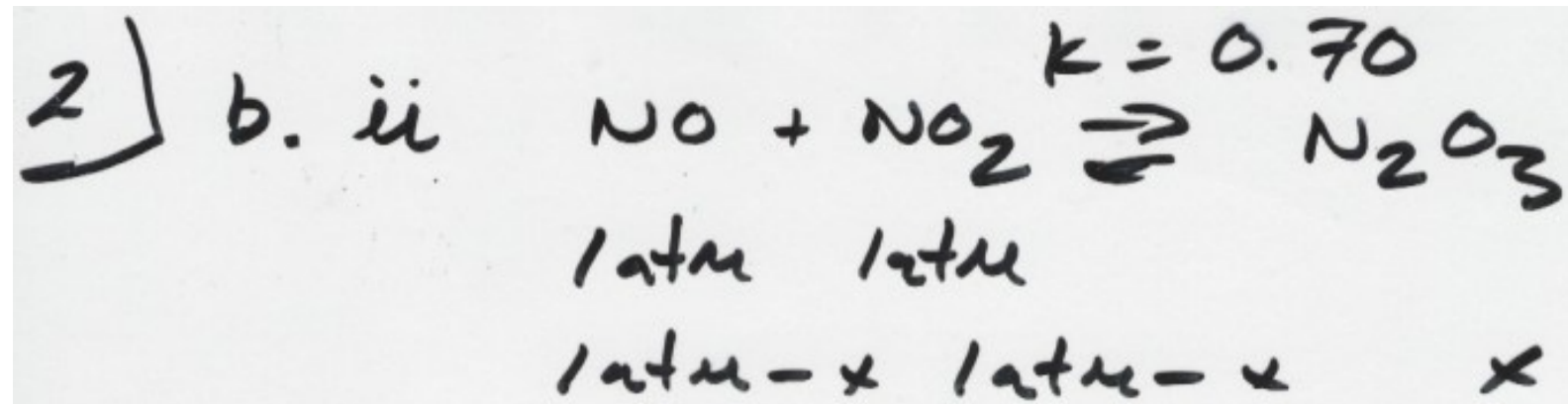g. $S_2O_3^{2-} + 4 \, OCl^- + 2OH^- \rightarrow$

$2 \, SO_4^{2-} + 4Cl^- + H_2O$

2) $2NO + O_2 \rightarrow 2NO_2$
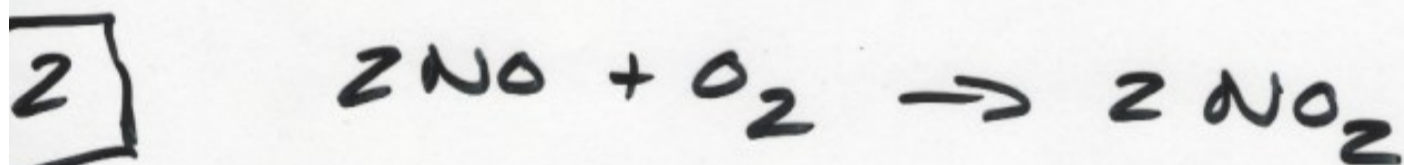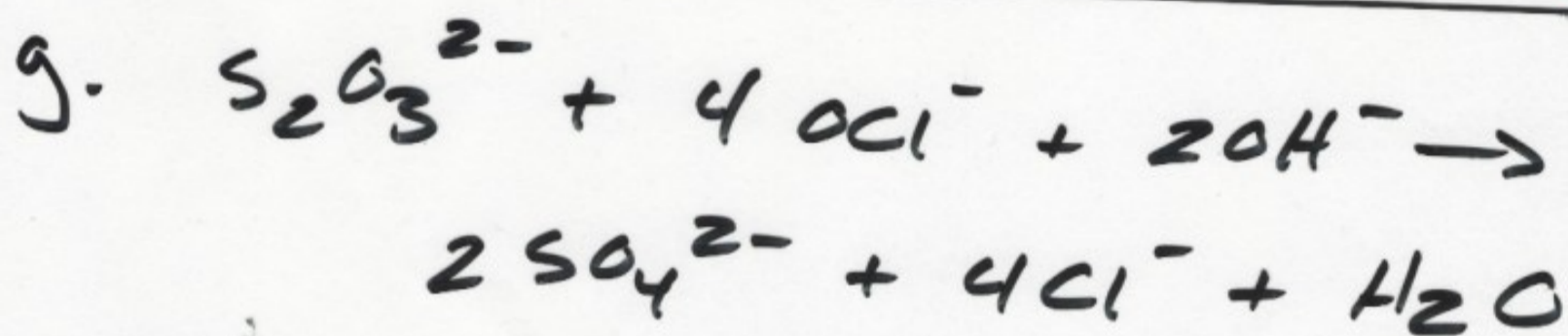
a.

$\rightarrow$
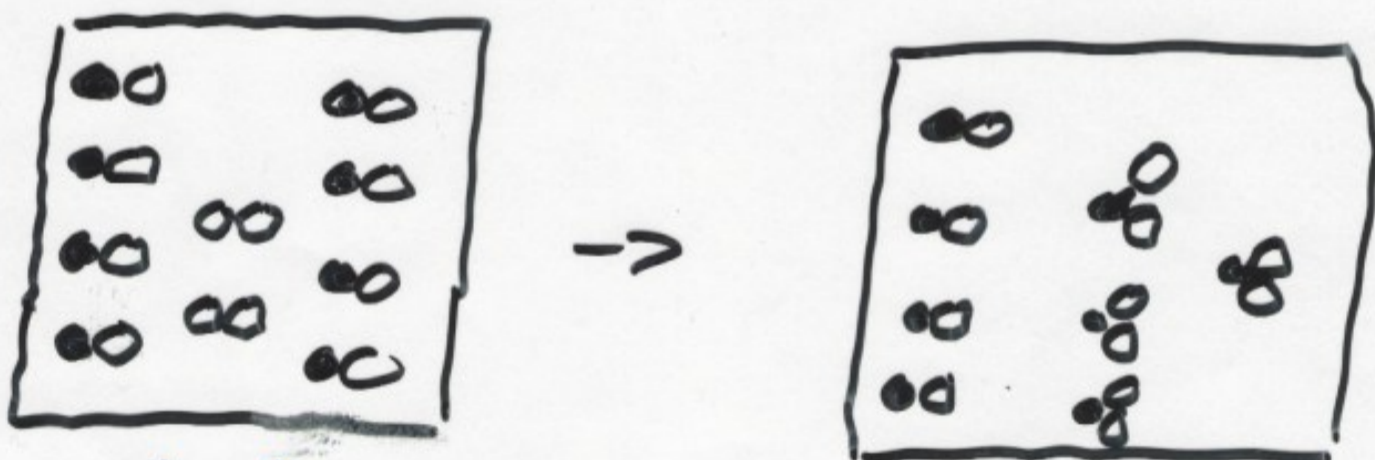
b. $\Delta G^\circ = -RT \ln k$

$0.87 \, kJ/mol = -\left(0.008314 \frac{kJ}{mol}\right)(298k) \ln k$

$-0.351 = \ln k$

$k = 0.70$

2) b. ii

$$NO + NO_2 \overset{k=0.70}{\rightleftharpoons} N_2O_3$$
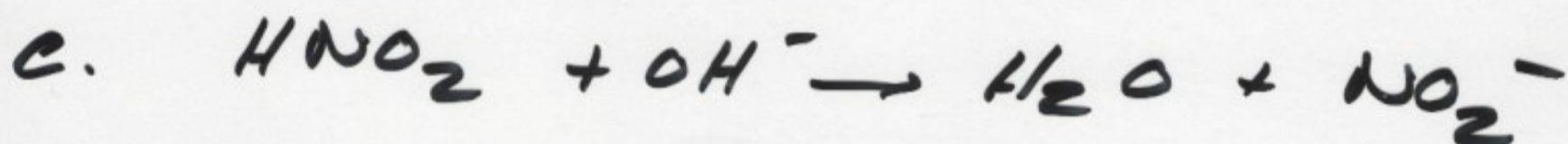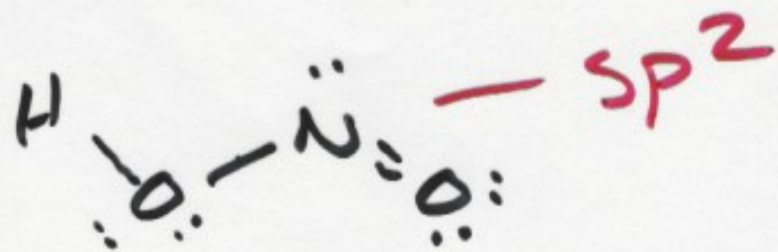
1 atm   1 atm

1 atm - x   1 atm - x   x
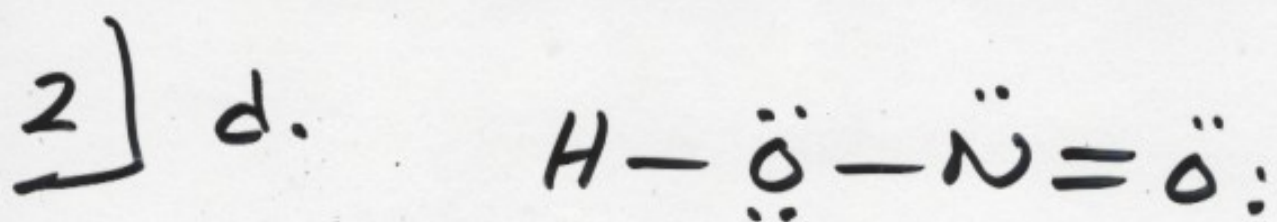
At equilibrium the the pressure of N_2O_3 will be less then 1 atm According to the kee ($k_{eq} = 0.70$) this is a nonspontaneous reaction and is reactant favored.

c. $$NO + NO_2 \rightleftharpoons N_2O_3 + 40kJ$$ ↓ Heat
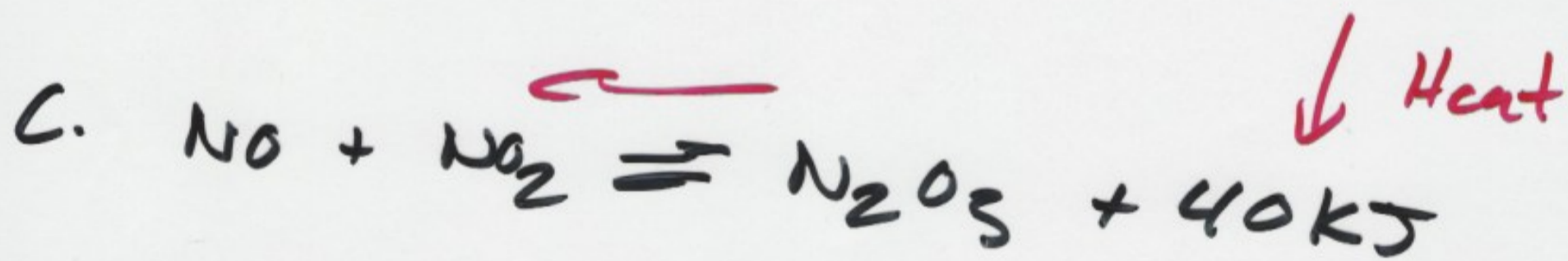
Disagree, exothermic reactions are less spontaneous (product favored) at higher temperatures.

2) d.

$$H - \ddot{O} - \ddot{N} = \ddot{O}:$$

$$H_{\diagdown} \ddot{O} - \ddot{N} = \ddot{O}: \quad - sp^2$$
$$\quad\quad :\ddot{O}:$$

c. $HNO_2 + OH^- \longrightarrow H_2O + NO_2^-$

$$20 \, mL \times \frac{0.100 \, mmol \, OH^-}{1 \, mL} \times \frac{1 \, mmol \, HNO_2}{1 \, mmol \, OH}$$

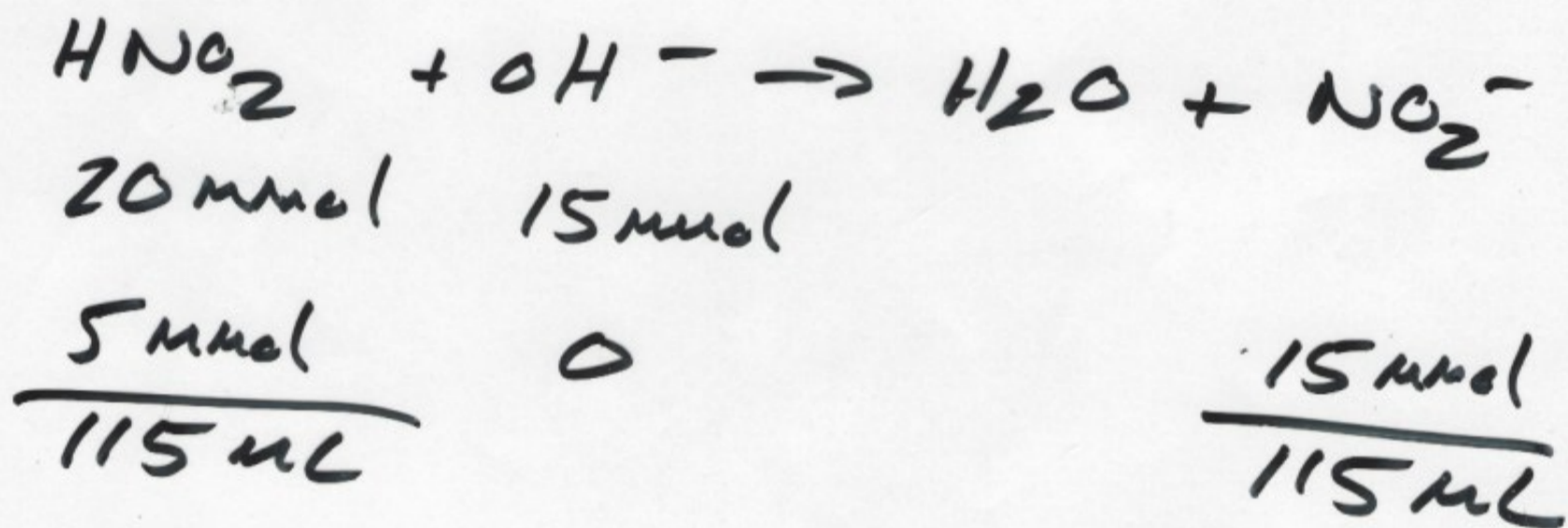$$\frac{2.0 \, mmol}{100 \, mL} = 2.0 \times 10^{-2} \, M \, HNO_2$$

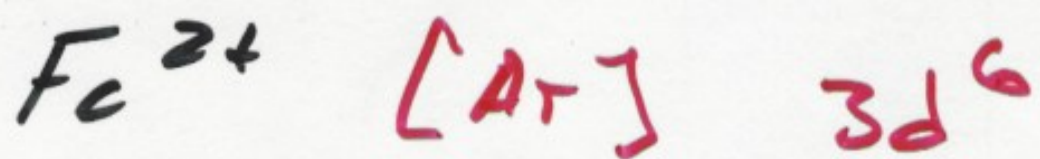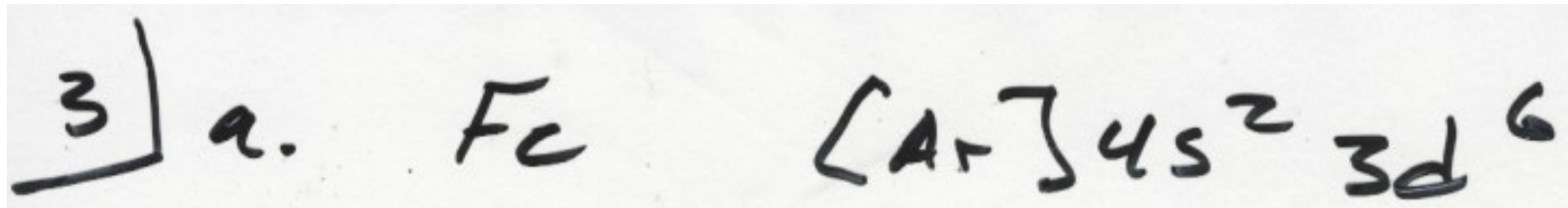ii. Halfway to the equivalence point the pH is 3.5

$$pH = pk_a + log \frac{[C.\,Base]}{[Acid]}$$

$$3.5 = pk_a + log(1)$$

$$pk_a = 3.5$$

f) NO_2^- would have a higher concentration. At 15 mL of 0.100 M KOH we are past the maximum buffer (half way to the equivalence pt.) and would have a higher concentration of NO_2^- then HNO_2

$$HNO_2 + OH^- \rightarrow H_2O + NO_2^-$$

20 mmol   15 mmol

$\dfrac{5\ mmol}{115\ mL}$ 0 $\dfrac{15\ mmol}{115\ mL}$

3) a. Fe $[Ar]4s^2 3d^6$

Fe^{2+} $[Ar]$ $3d^6$

b. Fe^{2+} has more electrons in its 3d orbitals then Fe^{3+} ($[Ar]3d^5$). This leads to more electron - electron repulsion and a bigger cloud.

c. Fe^{3+} is more highly charged and pulls harder on electrons then Fe^{2+}. This leads to stronger dipole - ion interactions with water.

3) d. $Fe^{2+} \rightarrow Fe^{3+} + 1e^-$

e. The 25 mL volumetric flask is good for making solutions. When titrating we would use the 25 mL buret. The buret allows us to add incremental volumes of solution

G. $7.531g \; Fe_2O_3 \times \dfrac{1 mol \; Fe_2O_3}{159.88g}$

$\times \dfrac{2 mol \; Fe}{1 mol \; Fe_2O_3} = 0.09421 \; mol \; Fe$

H. $7.531g \; Fe_2O_3 \times \dfrac{2(55.85g \; Fe)}{159.88g \; Fe_2O_3}$

$= \dfrac{5.267g \; Fe}{6.724g \; Fe} \times 100 = 78.33\%$

impurity

Fe

3] I.   lower %

FeO has a High % of iron by mass then Fe_2O_3. Assuming that the sample is all Fe_2O_3 gives a lower % of Fe then there actually is.

4] a. CS_2 has a higher boiling point due to being more massive and having stronger dispersion forces. Sulfur is a larger atom then Oxygen and more polarizable. COS is capable of dipole-dipole and dispersion forces (induced dipole - induced dipole)

4] b. $10.0 g \, CS_2 \times \dfrac{1 \, mol}{76.15 g} = 0.131 \, mol \, CS_2$

$P = \dfrac{nRT}{V}$

$$\dfrac{(0.131 \, mol \, CS_2)(0.08206 \, \frac{L \cdot atm}{mol \cdot k})(325 k)}{5.0 L}$$

$P = 0.70 \, atm$

5] a. HF is a weak acid and therefore has a low % ionization

b. $\dfrac{x}{0.0350} \times 100 = 13.0 \%$

$[H^+] = x = 4.55 \times 10^{-3}$

$HF + H_2O \rightleftharpoons F^- + H_3O^+$

$0.0350 - 4.55 \times 10^{-3} \quad 4.55 \times 10^{-3} \quad 4.55 \times 10^{-3}$

$k_a = \dfrac{(4.55 \times 10^{-3})^2}{(0.0350 - 4.55 \times 10^{-3})}$

$k_a = 6.80 \times 10^{-4}$

5 C. The more dilute the solution the greater the % ionization. $M_1 V_1 = M_2 V_2$

$$(0.035 M)(0.05 L) = (0.10 L) M_2$$

$$M_2 = 0.0175 M$$

$$HF + H_2O \rightleftharpoons H_3O^+ + F^-$$
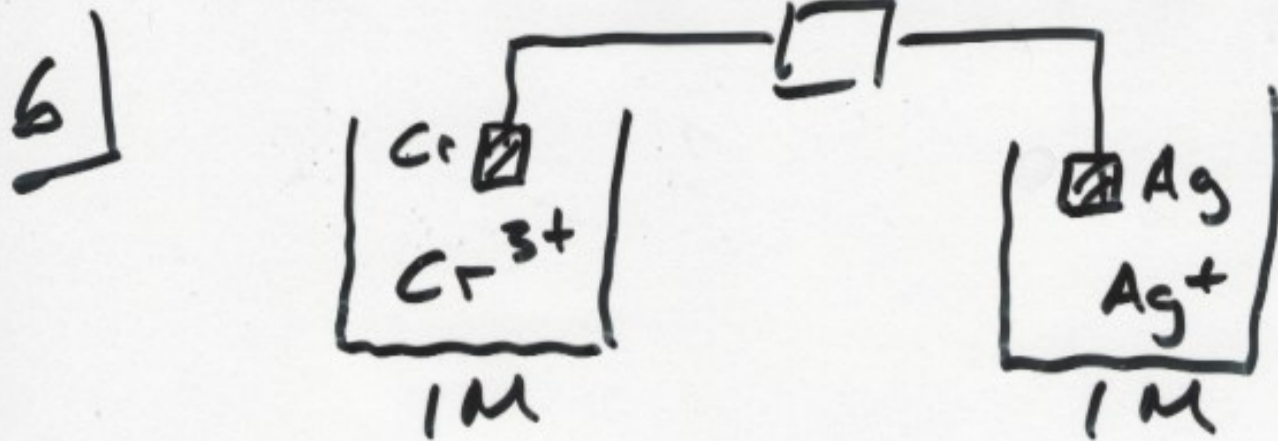
$$0.0175 - x \qquad\qquad x \qquad x$$

$$6.80 \times 10^{-4} = \frac{x^2}{0.0175 - x}$$

$$x = 3.45 \times 10^{-3}$$

% ionization $= \dfrac{3.45 \times 10^{-5}}{0.0175} \times 100$
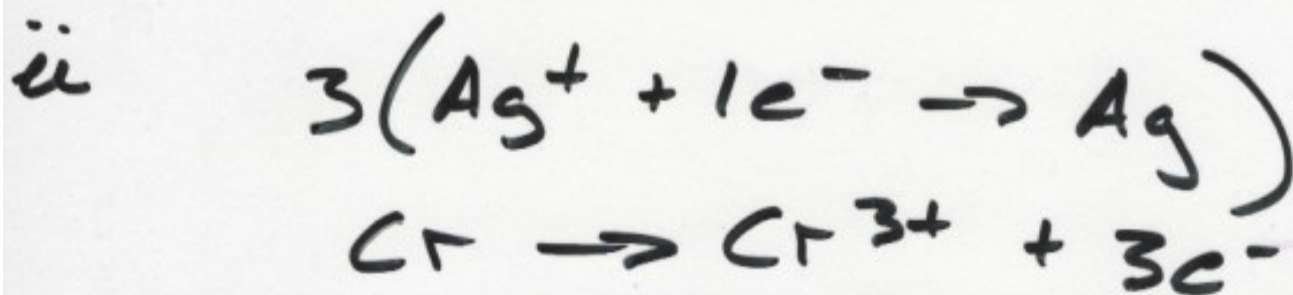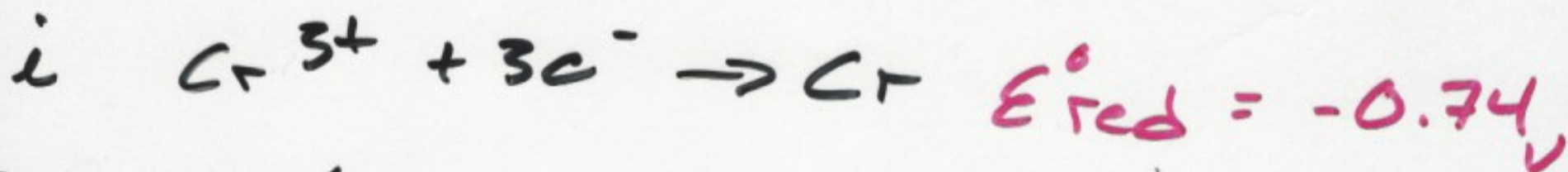
$$= 20. \%$$

$$13.0 \% < 20. \%$$

6)

a. A salt bridge is needed to balance the charge in the cell.

b. $Ag^+ + 1e^- \rightarrow Ag$ $\qquad$ E°_{red} $\quad +0.80v$

$Cr \rightarrow Cr^{3+} + 3e^-$ $\qquad$ E°_{ox} _____
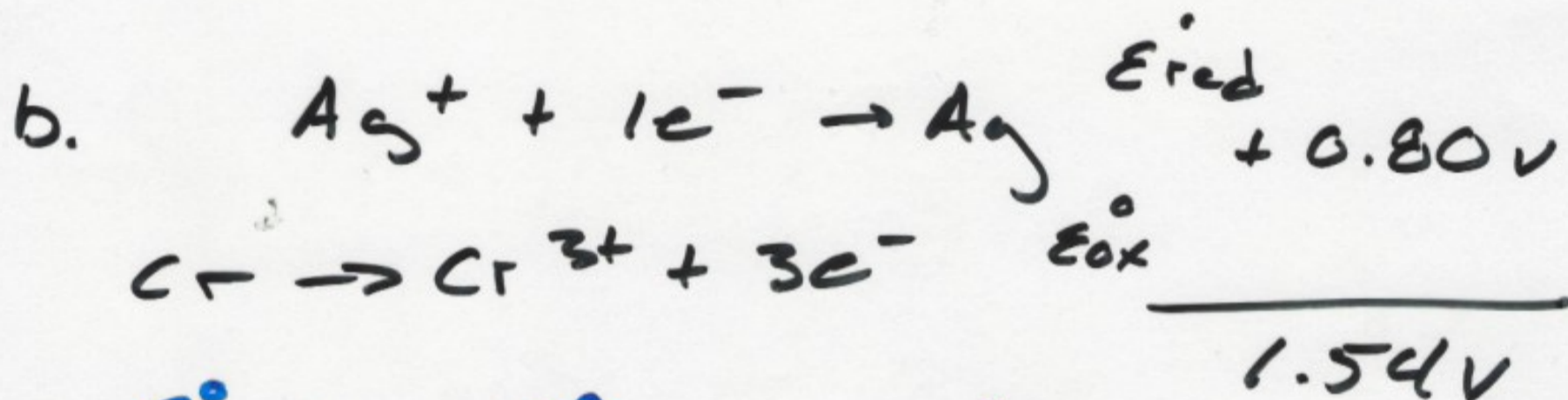
$\qquad\qquad\qquad\qquad\qquad\qquad\qquad$ $1.54v$

$E^\circ_{cell} - E^\circ_{red} = E^\circ_{ox}$

$1.54 - 0.80 = +0.74$

i $Cr^{3+} + 3e^- \rightarrow Cr$ $\quad E^\circ_{red} = -0.74v$

ii $\quad 3(Ag^+ + 1e^- \rightarrow Ag)$

$\qquad Cr \rightarrow Cr^{3+} + 3e^-$

_____

$3Ag^+ + Cr \rightarrow Cr^{3+} + 3Ag$

6] b iii $\quad \Delta G° = -n F E°$

$$= \frac{-3\ \text{mol}\ e^-}{\text{mol rxn}} \times \frac{96485\ c}{1\ \text{mol}\ e^-} \times \frac{1.54\ J}{c}$$

$-4.46 \times 10^5\ J/\text{mol rxn}$

7] a. N

b. $t_{1/2} = \dfrac{\ln 2}{k}$

$\quad k = \dfrac{\ln 2}{10.} = 6.9 \times 10^{-2}\ 1/\text{min}$

c. $\ln[\ ] = -kt + \ln[\ ]_c$

$\ln(1) = -\left(6.9 \times 10^{-2}\ \dfrac{1}{\text{min}}\right)(t) + \ln(64)$

$\quad t = 60.\ \text{min}$