



# Université d'Ottawa · University of Ottawa

Faculté des science  
Chimie

Faculty of Science  
Chemistry

## CHM 1301

DATE: December 9, 2013  
9:30 am

Dr. Rashmi Venkateswaran

TIME: 3 hours

Page 1 of 19

FULL STUDENT NAME: (Print) \_\_\_\_\_

STUDENT NUMBER: \_\_\_\_\_

### INSTRUCTIONS:

1. Write your answers on the exam. Please **BOLD** or **HIGHLIGHT** your final answer.
2. You are allowed to use electronic calculators during the exam (TI 30X, TI 34, TI 36, Casio 290 or 300).
3. Equations and constants are given on pages 17 and 18. A periodic table of the elements is included (page 19).
4. **Use the correct number of significant figures at all times.**
5. Marks allotted to each question are shown in brackets. The exam is out of 100.
7. Use the back of sheets for rough work. Three extra (blank) pages (14, 15 and 16) have also been included.
8. You are advised to read through the exam paper before you begin to write. Answer first those questions that you can do right away, proceeding to more difficult questions thereafter. Allow about ten minutes near the end of the exam period to read quickly through your answers, checking for obvious errors. **Good luck!**

Quest.	1	2	3	4	5	6	7	8	9	10	
Marks	/10	/10	/10	/10	/10	/10	/12	/8	/10	/10	/100

- (10) 1. A microwave oven operating at  $1.22 \times 10^8 \text{ nm}$  is used to heat  $150.0 \text{ mL}$  of water (roughly the volume of a tea cup) from  $20.0^\circ \text{C}$  to  $95.0^\circ \text{C}$ . Calculate the number of photons needed if  $92.0\%$  of microwave energy is converted to the thermal energy of water. ( $d_{\text{water}} = 1.00 \text{ g/mL}$ ;  $c_{\text{water}} = 4.184 \text{ J/g}^\circ\text{C}$ )

$$E_T = mc\Delta T = (150.0 \text{ mL})(1.00 \frac{\text{g}}{\text{mL}})(4.184 \frac{\text{J}}{\text{g}^\circ\text{C}})(95.0^\circ - 20.0^\circ) = 4.71 \times 10^4 \text{ J}$$

$$E_{\text{photon}} = h\nu = \frac{hc}{\lambda} = \frac{(6.626 \times 10^{-34} \text{ Js})(3.00 \times 10^8 \frac{\text{m}}{\text{s}})}{1.22 \times 10^8 + 10^{-9} \text{ m}} = 1.63 \times 10^{-24} \frac{\text{J}}{\text{photon}}$$

[5]  $E_T = E_{\text{photon}} \cdot \# \text{ photons} \Rightarrow \# \text{ photons} = \frac{E_T}{E_{\text{photon}}} = \frac{4.71 \times 10^4 \text{ J}}{1.63 \times 10^{-24} \frac{\text{J}}{\text{photon}}} = 2.89 \times 10^{28} \text{ photons}$

With energy loss,  $\# \text{ photons} = 2.89 \times 10^{28} / 0.92 = 3.14 \times 10^{28} \text{ photons}$

- (b) In a photoelectric effect experiment, photons having an energy of  $361.3 \text{ kJ/mol}$  are absorbed by a metal and the maximum kinetic energy of the resulting electrons is  $\text{KE} = 162.6 \text{ kJ/mol}$ .

- i) Calculate the binding energy (work function) of ONE electron in the metal.  $\text{KE}/e^- = \frac{162.6 \times 10^3 \text{ J/mol}}{6.022 \times 10^{23} e^-/\text{mol}} = 2.70 \times 10^{-19} \text{ J}$

$$E_{\text{photon}} = \frac{361.3 \times 10^3 \text{ J/mol}}{6.022 \times 10^{23} \text{ photons/mol}} = 6.00 \times 10^{-19} \text{ J}$$

$$\text{KE} = h\nu - \text{BE} \Rightarrow \text{BE} = h\nu - \text{KE} = (6.00 \times 10^{-19} - 2.70 \times 10^{-19}) \text{ J} = 3.30 \times 10^{-19} \text{ J/e}^-$$

- ii) Calculate the wavelength of the light.

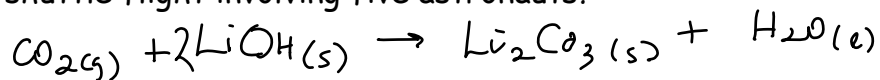
[5]  $E = h\nu = \frac{hc}{\lambda} \Rightarrow \lambda = \frac{hc}{E} = \frac{(6.626 \times 10^{-34} \text{ Js})(3.00 \times 10^8 \frac{\text{m}}{\text{s}})}{6.00 \times 10^{-19} \text{ J}} = 3.31 \times 10^{-7} \text{ m} = 331 \text{ nm}$

- iii) Calculate the velocity of the electrons.

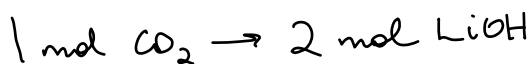
$$\text{KE} = \frac{1}{2} m v^2 \quad v^2 = \frac{2 \text{KE}}{m} = \frac{2(2.70 \times 10^{-19} \text{ J})}{9.11 \times 10^{-31} \text{ kg}}$$

$$v = 7.70 \times 10^5 \frac{\text{m}}{\text{s}}$$

- (10) 2. (a) The carbon dioxide exhaled by astronauts must be removed from the spacecraft atmosphere. One way to do this is to react the gaseous carbon dioxide with solid lithium hydroxide, which results in the formation of solid lithium carbonate and liquid water. The carbon dioxide output of an astronaut is about 1.0 kg/day. What is the minimum mass of lithium hydroxide required for a six-day space shuttle flight involving five astronauts?



$$n_{\text{CO}_2} = \frac{m}{\text{MM}} = \frac{1.0 \times 10^3 \text{ g}}{44.0 \frac{\text{g}}{\text{mol}}} = \left( 22.7 \frac{\text{mol}}{\text{person} \cdot \text{day}} \right) (6 \text{ day}) (5 \text{ person}) = 6.82 \times 10^2 \text{ mol CO}_2$$

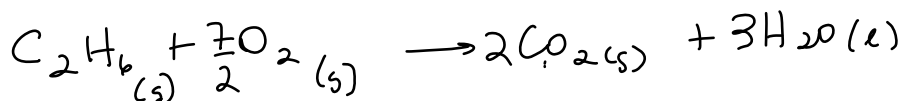


$$[5] \quad \therefore n_{\text{LiOH}} = 2n_{\text{CO}_2} = 2(6.82 \times 10^2 \text{ mol}) = 1.36 \times 10^3 \text{ mol}$$

$$m_{\text{LiOH}} = n \cdot \text{MM} = (1.36 \times 10^3 \text{ mol}) (6.941 + 16.0 + 1.00794 \frac{\text{g}}{\text{mol}}) \\ = 3.27 \times 10^4 \text{ g} = \boxed{32.7 \text{ kg}}$$

- (b) The combustion of what volume of ethane ( $\text{C}_2\text{H}_6$ ) measured at  $23.0^\circ\text{C}$  and 752 mm Hg would be required to heat 855 g of water from  $25.0^\circ\text{C}$  to  $98.0^\circ\text{C}$ ?  $c_{\text{water}} = 4.184 \text{ J/g}^\circ\text{C}$ ;  $\Delta_f H^\circ(\text{CO}_2) = -393.5 \text{ kJ/mol}$ ;  $\Delta_f H^\circ(\text{H}_2\text{O}(l)) = -285.8 \text{ kJ/mol}$ ;  $\Delta_f H^\circ(\text{C}_2\text{H}_6) = -84.7 \text{ kJ/mol}$

$$E_T = mc\Delta T = (855 \text{ g}) (4.184 \frac{\text{J}}{\text{g}^\circ\text{C}}) (98.0^\circ - 25.0^\circ\text{C}) \\ = 2.61 \times 10^5 \text{ J}$$



$$[5] \quad \Delta_r H^\circ = [2\Delta_f H^\circ_{\text{CO}_2} + 3\Delta_f H^\circ_{\text{H}_2\text{O}}] - [\Delta_f H^\circ_{\text{C}_2\text{H}_6}] \\ = [2(-393.5) + 3(-285.8) - (-84.7)] \frac{\text{kJ}}{\text{mol}}$$

$$= -1.5579 \times 10^6 \frac{\text{J}}{\text{mol}}$$

$$\rightarrow V = \frac{nRT}{P} = \frac{(0.168 \text{ mol}) (0.08314 \frac{\text{bar} \cdot \text{L}}{\text{mol} \cdot \text{K}}) (298.15 \text{ K})}{(\frac{752}{760} \times 1.01325 \text{ bar})} \\ = \boxed{4.11 \text{ L}}$$

$$1 \text{ mol C}_2\text{H}_6 \rightarrow 1.5597 \times 10^6 \text{ J/mol}$$

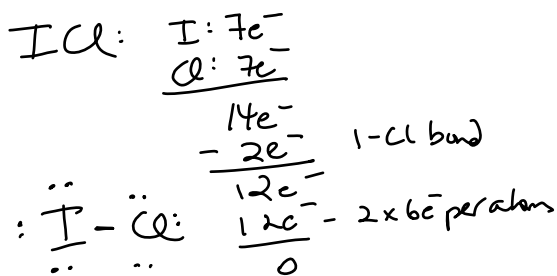
$$\times \rightarrow 2.61 \times 10^5 \text{ J}$$

$$1.5597 \times 10^6 \frac{\text{J}}{\text{mol}} \times = 2.61 \times 10^5 \text{ J}$$

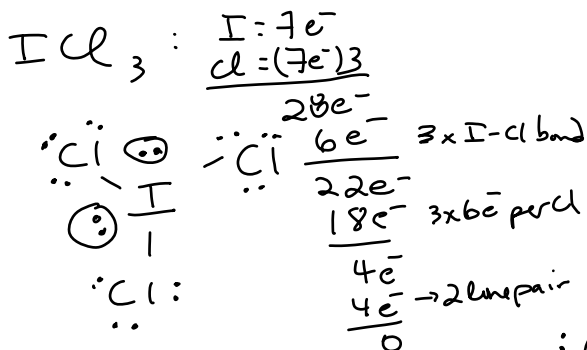
$$\times = 0.168 \text{ mol}$$

(10) 3. Iodine forms three compounds with chlorine:  $\text{ICl}$ ,  $\text{ICl}_3$ , and  $\text{ICl}_5$ .

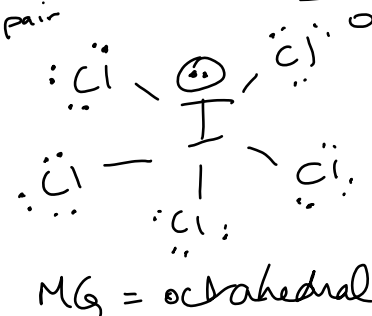
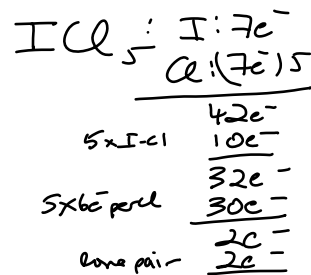
(a) Showing all your work, determine the molecular geometries of all three molecules.



[6]  $\text{MG} = \text{linear}$

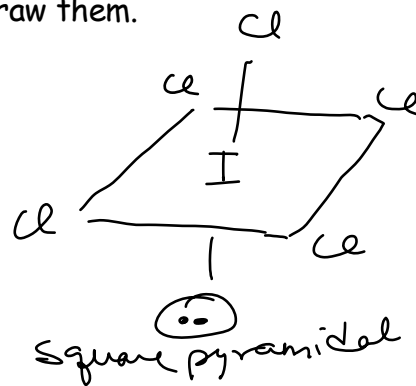
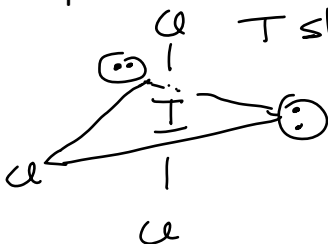


$\text{MG} = \text{trigonal bipyramidal}$



(b) What are the shapes of all three molecules? Draw them.

$\text{I-Cl} = \text{linear}$

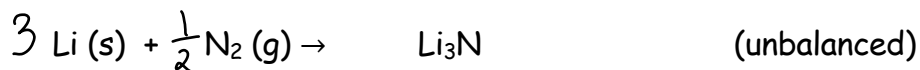


(c) What are the main angles in each of the three molecules? Are they polar?

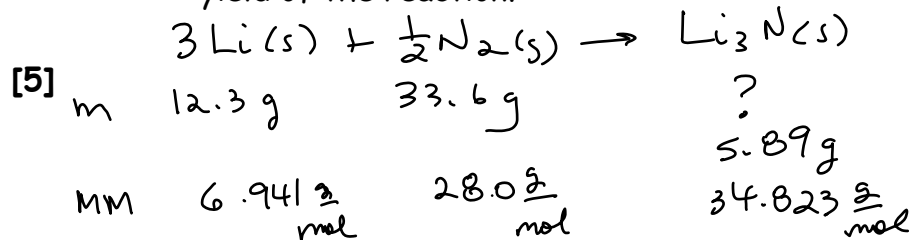
$\text{I-Cl} \rightarrow 180^\circ$  - polar because I & Cl different EN  
 $\text{I-Cl}_3 \rightarrow 180^\circ, 90^\circ$  - polar because of lone pairs vs Cl in plane  
 $\text{I-Cl}_5 \rightarrow 180^\circ, 90^\circ$  - polar because axial lone pair + Cl don't balance

[1]

(10) 4. (a) When heated, lithium reacts with nitrogen to form lithium nitride:



What is the theoretical yield of  $\text{Li}_3\text{N}$  when 12.3 g of Li is heated with 33.6 g of  $\text{N}_2$ ? If the actual yield of  $\text{Li}_3\text{N}$  is 5.89 g, what is the percent yield of the reaction?



$$n_{\text{Li}} = \frac{m}{\text{MM}} = \frac{12.3 \text{ g}}{6.941 \frac{\text{g}}{\text{mol}}} = 1.77 \text{ mol}$$

$$n_{\text{N}_2} = \frac{m}{\text{MM}} = \frac{33.6 \text{ g}}{28.02 \frac{\text{g}}{\text{mol}}} = 1.20 \text{ mol}$$

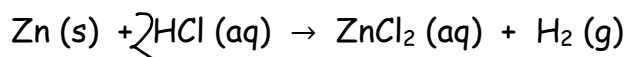
$\therefore$  Li is the limiting reagent (need 6 mol Li for each 1 mol  $\text{N}_2$ )

$$\therefore n_{\text{Li}_3\text{N}} \text{ formed} = \frac{1}{3} n_{\text{Li}} = \frac{1}{3} (1.77 \text{ mol}) = 0.591 \text{ mol}$$

$$m_{\text{Li}_3\text{N}} = (0.591 \text{ mol})(34.823 \frac{\text{g}}{\text{mol}}) = \boxed{20.6 \text{ g}}$$

$$\% \text{ yield} = \frac{\text{act}}{\text{theor}} \times 100\% = \frac{5.89 \text{ g}}{20.6 \text{ g}} \times 100\% = \boxed{28.6\%}$$

(b) A sample of zinc metal reacts completely with an excess of hydrochloric acid:



The hydrogen gas produced is collected over water at 25.0 °C. The volume of the gas is 7.80 L and the pressure is 0.980 atm. Calculate the mass of zinc metal consumed in the reaction.

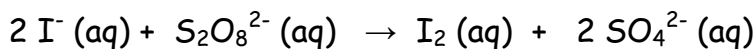
$$n = \frac{PV}{RT} = \frac{(0.980 \text{ atm})(7.80 \text{ L})}{(0.08206 \frac{\text{L}\cdot\text{atm}}{\text{mol}\cdot\text{K}})(298.2 \text{ K})} = 0.312 \text{ mol } \text{H}_2 \text{ collected}$$

[5]

$$n_{\text{H}_2} = n_{\text{Zn}} = 0.312 \text{ mol}$$

$$m_{\text{Zn}} = n \text{MM} = (0.312 \text{ mol})(65.38 \frac{\text{g}}{\text{mol}}) = \boxed{20.4 \text{ g}}$$

- (10) 5. (a) The following initial rate information was collected at 25°C for the reaction:



Experiment	$[\text{I}^-]_0$ (mol/L)	$[\text{S}_2\text{O}_8^{2-}]_0$ (mol/L)	Initial Rate (mol/L·s)
1	0.125	0.150	$4.4 \times 10^{-2}$
2	0.375	0.150	$1.3 \times 10^{-1}$
3	0.125	0.050	$1.5 \times 10^{-2}$

$$= k [\text{I}^-]^m [\text{S}_2\text{O}_8^{2-}]^n$$

- i) Determine the rate law expressed by this data.

$$\frac{\text{Rate}_2}{\text{Rate}_1} = \frac{k (0.375)^m (0.150)^n}{k (0.125)^m (0.150)^n} = \left(\frac{0.375}{0.125}\right)^m = 3^m = \frac{1.3}{4.4 \times 10^{-2}} = 3 \quad 3^m = 3 \quad m=1$$

$$\frac{\text{Rate}_1}{\text{Rate}_3} = \frac{k (0.125)^m (0.150)^n}{k (0.125)^m (0.050)^n} = \left(\frac{0.150}{0.050}\right)^n = 3^n = \frac{4.4 \times 10^{-2}}{1.5 \times 10^{-2}} = 3 \quad 3^n = 3 \quad n=1$$

$$\text{Rate} = k [\text{I}^-] [\text{S}_2\text{O}_8^{2-}]$$

- ii) Calculate the rate constant with the CORRECT units.

use any trial Trial 3  $1.5 \times 10^{-2} \frac{\text{mol}}{\text{L}\cdot\text{s}} = k \left(0.125 \frac{\text{mol}}{\text{L}}\right) \left(0.050 \frac{\text{mol}}{\text{L}}\right)$

$$k = 2.4 \frac{\text{L}}{\text{mol}\cdot\text{s}}$$

- b) The light-emitting decay of excited mercury atoms is first order, with a half-life of  $4.20 \times 10^{-7}$  s. A sample contains  $4.5 \times 10^{-6}$  mol/L of excited mercury atoms.

$$t_{1/2} = 4.20 \times 10^{-7} \text{ s} \quad k = \frac{\ln 2}{t_{1/2}} = \frac{\ln 2}{4.20 \times 10^{-7} \text{ s}} = 1.65 \times 10^6 \text{ s}^{-1}$$

- i) At what time will the concentration be  $4.5 \times 10^{-7}$  mol/L?

$$\ln \frac{[A]_t}{[A]_0} = -kt \quad \ln \left(\frac{4.5 \times 10^{-7} \frac{\text{mol}}{\text{L}}}{4.5 \times 10^{-6} \frac{\text{mol}}{\text{L}}}\right) = -(1.65 \times 10^6 \text{ s}^{-1}) t$$

$$t = 1.40 \times 10^{-6} \text{ s} = 1.40 \mu\text{s}$$

- ii) What will the concentration of mercury atoms be after  $2.5 \times 10^{-6}$  s?

$$\ln \frac{[A]_t}{[A]_0} = -kt \quad \ln \left(\frac{[A]_t}{4.5 \times 10^{-6} \frac{\text{mol}}{\text{L}}}\right) = -(1.65 \times 10^6 \text{ s}^{-1})(2.5 \times 10^{-6} \text{ s})$$

$$[A]_t = 7.3 \times 10^{-8} \frac{\text{mol}}{\text{L}}$$

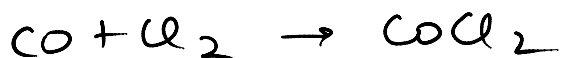
$$\left[ \begin{aligned} \ln \frac{[A]_t}{4.5 \times 10^{-6}} &= -4.125 \\ e^{-4.125} &= \frac{[A]_t}{4.5 \times 10^{-6}} \\ [A]_t &= 4.5 \times 10^{-6} e^{-4.125} \end{aligned} \right]$$

b) The reaction of CO with Cl<sub>2</sub> gives phosgene (COCl<sub>2</sub>), a nerve gas used in World War I. Even though the stoichiometry is simple, the mechanism has several steps:

1.  $\text{Cl}_2(\text{g}) \xrightleftharpoons[k_{-1}]{} 2 \text{Cl}(\text{g})$  Fast reversible equilibrium
2.  $\text{Cl}(\text{g}) + \text{CO}(\text{g}) \xrightarrow{k_2} \text{COCl}(\text{g})$  Slow
3.  $\text{COCl}(\text{g}) + \text{Cl}(\text{g}) \xrightarrow{k_3} \text{COCl}_2(\text{g})$  Fast

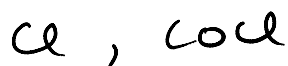
- i) Show that the elementary reactions of the proposed mechanism can be added to provide a balanced equation for the reaction.

[1]



- ii) Identify any intermediates in the mechanism.

[1]



- iii) What is the rate law yielded by this mechanism?

[2]

$$\text{Rate}_{\text{fwd}} \textcircled{1} = \text{Rate}_{\text{rev}} \textcircled{1} \quad \textcircled{A} \quad k_1 [\text{Cl}_2] = k_{-1} [\text{Cl}]^2$$

Rate is determined by slow step (RDS)  $\textcircled{2}$

$$\textcircled{B} \quad \therefore \text{Rate} = k_2 [\text{Cl}] [\text{CO}]$$

but  $[\text{Cl}]$  is an intermediate!

$$\text{using } \textcircled{A} \text{ solve for } [\text{Cl}] \Rightarrow [\text{Cl}] = \sqrt{\frac{k_1}{k_{-1}} [\text{Cl}_2]} = \left(\frac{k_1}{k_{-1}}\right)^{1/2} [\text{Cl}_2]^{1/2}$$

$$\text{from } \textcircled{B} \quad \text{Rate} = k_2 [\text{Cl}] [\text{CO}]$$

$$= k_2 \left(\frac{k_1}{k_{-1}}\right)^{1/2} [\text{Cl}_2]^{1/2} [\text{CO}]$$

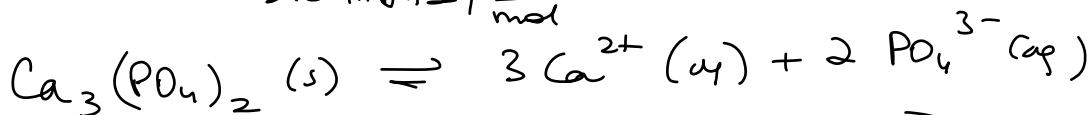
$$= k' [\text{Cl}_2]^{1/2} [\text{CO}], \quad \text{where } k' = k_2 \left(\frac{k_1}{k_{-1}}\right)^{1/2}$$

(10) 6. (a) The solubility of calcium phosphate in water is  $3.5 \times 10^{-5}$  g/L. Use this information to calculate  $K_{sp}$  for the salt.

$$\text{Ca}_3(\text{PO}_4)_2 \Rightarrow \text{MM} = 3(40.078 \frac{\text{g}}{\text{mol}}) + 2(30.973762 \frac{\text{g}}{\text{mol}}) + 8(15.9994 \frac{\text{g}}{\text{mol}})$$

$$= 310.176724 \frac{\text{g}}{\text{mol}}$$

$$s = \frac{[5] \text{ solubility}}{\text{MM}} (\text{Ca}_3(\text{PO}_4)_2) = \frac{(3.5 \times 10^{-5} \frac{\text{g}}{\text{L}})}{310.176724 \frac{\text{g}}{\text{mol}}} = 1.1 \times 10^{-7} \frac{\text{mol}}{\text{L}}$$



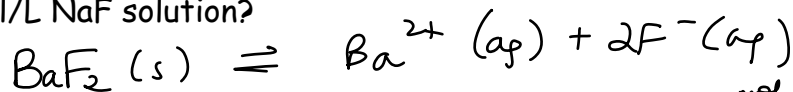
I	some	—	2s
C	-s	3s	2s
E	some	3s	2s

$$K_{sp} = [\text{Ca}^{2+}]^3 [\text{PO}_4^{3-}]^2 = (3s)^3 (2s)^2 = 27s^3 \cdot 4s^2 = 108 s^5$$

$$= 108 (1.1 \times 10^{-7})^5$$

$$= \underline{\underline{2.0 \times 10^{-33}}}$$

(b) What mass of  $\text{BaF}_2$  ( $K_{sp} = 1.8 \times 10^{-7}$ ) will dissolve in 0.500 L of 0.100 mol/L NaF solution?



[5]	I	some	—	0.100 $\frac{\text{mol}}{\text{L}}$
	C	-s	s	2s
	E	some	s	2s + 0.100

$$K_{sp} = 1.8 \times 10^{-7} = [\text{Ba}^{2+}][\text{F}^-]^2 = s(2s + 0.100)^2 = 1.8 \times 10^{-7}$$

Assume  $2s \ll 0.100 \therefore 2s + 0.100 \approx 0.100$

$$s(0.100)^2 = 1.8 \times 10^{-7}$$

$$s = 1.8 \times 10^{-5} \frac{\text{mol}}{\text{L}}$$

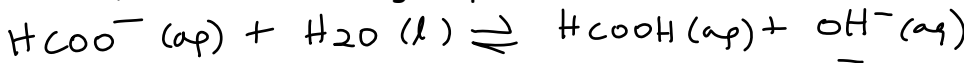
$$n_{\text{BaF}_2} = CV = (1.8 \times 10^{-5} \frac{\text{mol}}{\text{L}})(0.500 \text{ L}) = 9.0 \times 10^{-6} \text{ mol}$$

$$m_{\text{BaF}_2} = n \text{MM} = (9.0 \times 10^{-6} \text{ mol})(137.327 + 2(18.998403) \frac{\text{g}}{\text{mol}})$$

$$= 1.6 \times 10^{-3} \text{ g} = \boxed{1.6 \text{ mg}}$$

(12) 7. (a) Consider the titration of 30.0 mL of 0.200 mol/L HCOONa (aq) with 6.0 mol/L HCl (aq). ( $K_a$  for HCOOH is  $1.8 \times 10^{-4}$ )

i) What is the original pH of the solution of HCOONa?  $\text{HCOONa} \rightarrow \text{Na}^+ + \text{HCOO}^-$  (conj. base)



I	0.200 mol/L
C	-x
E	0.200 - x

-	-
x	x
x	x

$$K_b = \frac{K_w}{K_a} = \frac{1.0 \times 10^{-14}}{1.8 \times 10^{-4}} = 5.6 \times 10^{-11} = \frac{[\text{HCOOH}][\text{OH}^-]}{[\text{HCOO}^-]} = \frac{x^2}{0.200 - x} = 5.6 \times 10^{-11}$$

assume  $x \ll 0.200 \therefore 0.200 - x \approx 0.200 \therefore \frac{x^2}{0.200} = 5.6 \times 10^{-11} \quad x = 3.3 \times 10^{-6} \text{ mol/L} = [\text{OH}^-]$

$\text{pOH} = -\log[\text{OH}^-] = -\log(3.3 \times 10^{-6}) = 5.48 \quad \text{pH} = 14.00 - \text{pOH} = 14.00 - 5.48$

ii) What is the pH of the solution halfway through the titration? = 8.52

First find equivalence pt!  $C_A V_A = C_B V_B$  (since 1:1)

$$(6.0 \frac{\text{mol}}{\text{L}}) V_A = (0.200 \frac{\text{mol}}{\text{L}})(30.0 \text{ mL}) \quad V_A = 1.00 \text{ mL}$$

[6]  $\therefore$  The  $V_{\text{Aeq}}$  halfway through titration is 0.50 mL.

$\text{HCl} (\text{aq}) + \text{HCOO}^- (\text{aq}) \rightarrow \text{HCOOH} (\text{aq}) + \text{Cl}^- (\text{aq})$			
$6.0 \frac{\text{mol}}{\text{L}} (\frac{0.50 \text{ mL}}{30.50 \text{ mL}})$	$0.200 \frac{\text{mol}}{\text{L}} (\frac{30.00 \text{ mL}}{30.50 \text{ mL}})$	-	-
I	0.098 mol/L	0.197 mol/L	0.098
C	-0.098	-0.098	0.098
E	0	0.099	0.098

Left with Buffer!

$\text{pH} = \text{p}K_a + \log \frac{[\text{A}^-]}{[\text{HA}]}$

$\text{pH} = -\log(1.8 \times 10^{-4}) + \log(\frac{0.099}{0.098})$

= 3.75

iii) What is the pH at the equivalence point?

$\text{HCl} (\text{aq}) + \text{HCOO}^- (\text{aq}) \rightarrow \text{HCOOH} (\text{aq}) + \text{Cl}^- (\text{aq})$			
$6.0 \frac{\text{mol}}{\text{L}} (\frac{1.00 \text{ mL}}{31.00 \text{ mL}})$	$0.200 \frac{\text{mol}}{\text{L}} (\frac{30.00 \text{ mL}}{31.00 \text{ mL}})$	-	-
I	0.1935 mol/L	0.1935	0.1935
C	-0.1935	-0.1935	0.1935
E	0	0.1935	0.1935

must RE DO ICE!

$\text{HCOOH} (\text{aq}) + \text{H}_2\text{O} (\text{l}) \rightleftharpoons \text{HCOO}^- (\text{aq}) + \text{H}_3\text{O}^+ (\text{aq})$			
I	0.1935	-	-
C	-x	x	x
E	0.1935 - x	x	x

$$K_a = \frac{[\text{HCOO}^-][\text{H}_3\text{O}^+]}{[\text{HCOOH}]} = \frac{x^2}{0.1935 - x} = 1.8 \times 10^{-4} \Rightarrow x^2 = 1.8 \times 10^{-4} (0.1935 - x)$$

$$x^2 + 1.8 \times 10^{-4} x - 3.5 \times 10^{-5} = 0 \quad x = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a}$$

$\text{pH} = -\log[\text{H}^+] = -\log(5.8 \times 10^{-3}) = 2.23$   
 or 2.24

(use  $\pm$  root)  $= \frac{-1.8 \times 10^{-4} \pm \sqrt{(1.8 \times 10^{-4})^2 - 4(1)(-3.5 \times 10^{-5})}}{2}$   
 $= 5.8 \times 10^{-3} \frac{\text{mol}}{\text{L}} = [\text{H}^+]$

HCl reacts with  $\text{HCOO}^-$   $\rightarrow$  not  $\rightleftharpoons$  because of strong acid

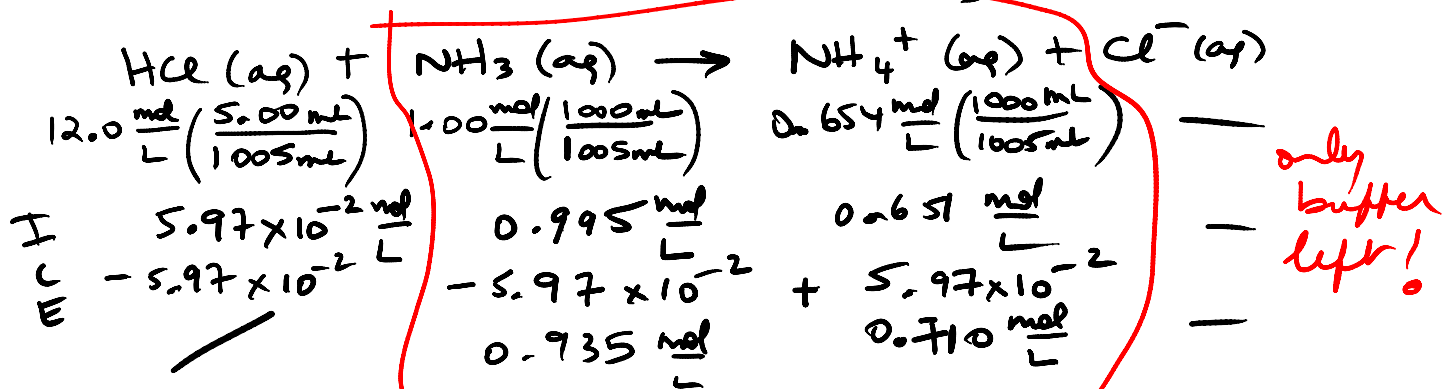
$$K_b = 1.8 \times 10^{-5}$$

(b) What is the pH of a buffer solution containing 35.0 g of  $\text{NH}_4\text{Cl}$  dissolved in 1.00 L of 1.00 mol/L  $\text{NH}_3$ ?  $\text{MM}_{\text{NH}_4\text{Cl}} = 14.01 + 4(1.01) + 35.45 \frac{\text{g}}{\text{mol}} = 53.5 \frac{\text{g}}{\text{mol}}$

$$n_{\text{NH}_4\text{Cl}} = \frac{35.0 \text{ g}}{53.5 \frac{\text{g}}{\text{mol}}} = 0.654 \text{ mol in } 1.00 \text{ L} \Rightarrow 0.654 \frac{\text{mol}}{\text{L}} \text{ of } \text{NH}_4^+$$

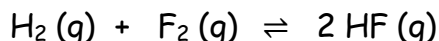
$$\begin{aligned} [6] \quad \text{pH} &= \text{p}K_a + \log \frac{[\text{A}^-]}{[\text{HA}]} \\ &= -\log \left( \frac{K_w}{K_b} \right) + \log \frac{[\text{A}^-]}{[\text{HA}]} \\ &= -\log \left( \frac{1.00 \times 10^{-14}}{1.8 \times 10^{-5}} \right) + \log \left( \frac{1.00 \frac{\text{mol}}{\text{L}}}{0.654 \frac{\text{mol}}{\text{L}}} \right) \\ &= \underline{9.44} \end{aligned}$$

ii) Calculate the pH of the system if 5.0 mL of 12.0 mol/L HCl is added to the buffer. *HCl added reacts with  $\text{NH}_3$*



$$\begin{aligned} \text{pH} &= \text{p}K_a + \log \frac{[\text{A}^-]}{[\text{HA}]} = -\log 5.6 \times 10^{-10} + \log \left( \frac{0.935}{0.710} \right) \\ &= 9.37 \quad (\text{reasonable} \rightarrow \text{pH decreases when we add acid but only slightly}) \end{aligned}$$

(8) 8. Hydrogen fluoride is a highly reactive gas. It has many industrial uses, but the most familiar property of HF is its ability to react with glass. As a result, HF is used to etch glass and frost the inner surfaces of light bulbs. Hydrogen fluoride gas must be stored in stainless steel containers and aqueous solutions must be stored in plastic bottles. Hydrogen fluoride can be produced  $H_2(g)$ , and  $F_2(g)$ :



In a particular experiment, 3.00 bar of hydrogen gas and 2.00 bar of fluorine gas are added to a 1.50 L flask at 350 K. Calculate the equilibrium partial pressures of all species.

(8) 8. Hydrogen fluoride is a highly reactive gas. It has many industrial uses, but the most familiar property of HF is its ability to react with glass. As a result, HF is used to etch glass and frost the inner surfaces of light bulbs. Hydrogen fluoride gas must be stored in stainless steel containers and aqueous solutions must be stored in plastic bottles. Hydrogen fluoride can be produced  $H_2(g)$ , and  $F_2(g)$ :

*\* sqrt  
# = 101.2*

*$P_{HF} = 2x = 3.65 \text{ bar}$*

*$P_{H_2} = 3.00 - x = 1.18 \text{ bar}$*

*$P_{F_2} = 2.00 - x = 0.18 \text{ bar}$*

$H_2(g) + F_2(g) \rightleftharpoons 2 HF(g) \quad K = 64.0$

I      3.00 bar    2.00 bar    —

In a particular experiment, 3.00 bar of hydrogen gas and 2.00 bar of fluorine gas are added to a 1.50 L flask at 350 K. Calculate the equilibrium partial pressures of all species.

C      -x      -x      2x

E      3-x      2-x      2x

$K = \frac{P_{HF}^2}{P_{H_2} P_{F_2}} = \frac{(2x)^2}{(3-x)(2-x)} = 64$

$4x^2 = 384 - 320x + 64x^2$

$60x^2 - 320x + 384 = 0$

$x = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a} = \frac{320 \pm \sqrt{(-320)^2 - 4(60)(384)}}{120} = 1.52 \text{ bar}$

$\frac{4x^2}{6-5x+x^2} = \frac{64}{1}$

[5] A coin dealer, offered a rare silver coin, suspected that it might be a counterfeit nickel copy. The dealer heated the coin, of mass 15.5 g, to 100.0 °C in boiling water and then dropped the hot coin into 21.5 g of water in a coffee cup calorimeter at 15.5 °C. The temperature of the water rose to 21.5 °C. Was the coin made of silver or nickel?

$c_{Ag} = 0.240 \text{ J/g}^\circ\text{C}$        $c_{Ni} = 0.440 \text{ J/g}^\circ\text{C}$

(c) A coin dealer, offered a rare silver coin, suspected that it might be a counterfeit nickel copy. The dealer heated the coin, of mass 15.5 g, to 100.0 °C in boiling water and then dropped the hot coin into 21.5 g of water in a coffee cup calorimeter at 15.5 °C. The temperature of the water rose to 21.5 °C. Was the coin made of silver or nickel?

$c_{Ag} = 0.240 \text{ J/g}^\circ\text{C}$        $c_{Ni} = 0.440 \text{ J/g}^\circ\text{C}$

$q_{H_2O} = -q_{met}$

$(m c \Delta T)_{H_2O} = -(m c \Delta T)_{met}$

[3]  $(21.5g)(4.184 \frac{J}{g^\circ C})(21.5^\circ - 15.5^\circ) = -(15.5g)(c)(21.5 - 100.0^\circ)$

$c = 0.443 \frac{J}{g^\circ C} \Rightarrow \text{It's Nickel!}$

[3]

(10) 9. Police officers confiscate a packet of white powder that they believe contains heroin. Purification by a forensic chemist yields a 38.70 mg sample for combustion analysis. This sample gives 97.46 mg of carbon dioxide and 20.81 mg of water. A second sample is analysed for its nitrogen content, which is found to be 3.8%. Show by calculations whether these data are consistent with the formula for heroin,  $C_{21}H_{22}NO_5$ .

$$n_C = n_{CO_2} = \frac{m}{MM} = \frac{97.46 \times 10^{-3} \text{ g}}{44.01 \text{ g/mol}} = 2.214 \times 10^{-3} \text{ mol}$$

$$m_C = n_C MM_C = (2.214 \times 10^{-3} \text{ mol})(12.01 \frac{\text{g}}{\text{mol}}) = 2.660 \times 10^{-2} \text{ g}$$

$$n_{H_2O} = \frac{m}{MM} = \frac{20.81 \times 10^{-3} \text{ g}}{18.02 \text{ g/mol}} = 1.155 \times 10^{-3} \text{ mol}$$

$$n_H = 2n_{H_2O} = 2(1.155 \times 10^{-3} \text{ mol}) = 2.310 \times 10^{-3} \text{ mol}$$

$$m_H = n_H MM_H = (2.310 \times 10^{-3} \text{ mol})(1.01 \frac{\text{g}}{\text{mol}}) = 2.333 \times 10^{-3} \text{ g}$$

$$m_N = 3.8\% (38.70 \text{ mg}) = 1.5 \text{ mg}$$

$$n_N = \frac{m}{MM} = \frac{1.471 \times 10^{-3} \text{ g}}{14.01 \text{ g/mol}} = 1.05 \times 10^{-4} \text{ mol}$$

$$m_O = m_{\text{comp}} - m_C - m_H - m_N = 38.70 \text{ mg} - 26.60 \text{ mg} - 2.333 \text{ mg} - 1.5 \text{ mg} = 8.3 \text{ mg}$$

$$\therefore n_O = \frac{m}{MM} = \frac{8.3 \times 10^{-3} \text{ g}}{16.01 \text{ g/mol}} = 5.2 \times 10^{-4} \text{ mol}$$

$$\begin{array}{cccc} C & H & N & O \\ \frac{2.214 \times 10^{-3}}{1.05 \times 10^{-4}} & \frac{2.310 \times 10^{-3}}{1.05 \times 10^{-4}} & \frac{1.05 \times 10^{-4}}{1.05 \times 10^{-4}} & \frac{5.2 \times 10^{-4}}{1.05 \times 10^{-4}} \end{array}$$



Yes it is cocaine

(10) 10. Your parents have offered to pay your tuition for the fall semester of University if you can explain to them why you have selected the courses you chose and why they will be potentially helpful in getting you a job. You, of course, choose Chemistry to demonstrate how important it is to our lives today. Using any one of the topics covered during the semester, write a letter of minimum length 1 page and using at least 3 chemical concepts within your chosen topic (clearly explained and not just repeated from class) to demonstrate why a knowledge of chemistry is essential in our daily lives.

Various answers!







## Equations and Constants

### Gas Laws

$$PV = nRT$$

$$P_{\text{Total}} = P_1 + P_2 + P_3 + \dots$$

$$d = m/V = P(\text{MM}) / RT$$

$$KE = (1/2)mv_{\text{av}}^2$$

$$u_{\text{rms}} = \sqrt{\frac{3RT}{M}}$$

$$\frac{\text{Rate}_A}{\text{Rate}_B} = \sqrt{\frac{M_b}{M_a}}$$

$$P + \frac{n^2 a}{V^2} (V - nb) = nRT$$

### Equilibrium

$$K_p = K(RT)^{\Delta n}$$

$$K = \frac{[C]^c [D]^d}{[A]^a [B]^b}$$

$$K_{\text{sp}} = [C]^c [D]^d$$

### Acid/Base

$$\text{pOH} = -\log [\text{OH}^-]$$

$$\text{pH} = -\log [\text{H}^+]$$

$$\text{pH} + \text{pOH} = 14$$

$$K_a \cdot K_b = K_w$$

$$\text{pH} = \text{p}K_a + \log \frac{[\text{A}^-]}{[\text{HA}]}$$

$$\text{pH} = \frac{\text{p}K_{a1} + \text{p}K_{a2}}{2}$$

$$C_1 V_1 = C_2 V_2$$

$$\ln \left( \frac{K_1}{K_2} \right) = \frac{\Delta H^\circ}{R} \left( \frac{1}{T_2} - \frac{1}{T_1} \right)$$

### Electrochemistry

$$\Delta G^\circ = -nFE^\circ$$

$$E = E^\circ - \frac{RT}{nF} \ln(Q)$$

$$\text{Or } E = E^\circ - \frac{0.059}{n} \log(Q) \quad \text{at } 25^\circ\text{C}$$

$$Q = I \cdot t$$

### Quantum Mechanics

$$\lambda \cdot \nu = c$$

$$E = h \cdot \nu$$

$$m = \frac{h}{\lambda c}$$

$$E = \frac{1}{2} m u^2$$

$$\lambda = h/mu$$

$$\text{Energy of state} = -2.178 \times 10^{-18} \text{ J} / n^2$$

$$\Delta x \cdot \Delta p \geq h / 4\pi$$

$$E = -C(1/n^2)$$

### Liquids and Colligative Properties

$$\ln(P_1/P_2) = \Delta H^\circ / R (1/T_2 - 1/T_1)$$

$$P_{\text{solution}} = X_{\text{solvent}} \cdot P_{\text{solvent}}^\circ$$

$$\Delta T_{\text{BP}} = K_{\text{BP}} \cdot m$$

$$\Delta T_{\text{FP}} = K_{\text{FP}} \cdot m$$

$$\Delta T = K \cdot m \cdot i$$

$$\Pi = cRT$$

$$\text{molality} = n(\text{solute})/m(\text{solvent (kg)})$$

### General INFORMATION

$$n = m/\text{MM}$$

$$d = m/V$$

$$C = n/V$$

$$1 \text{ atm} = 760 \text{ Torr} = 760 \text{ mm Hg} = 1.01325 \times 10^5 \text{ Pa} = 1.01325 \text{ bar} = 101.325 \text{ kPa}$$

Thermodynamics	Kinetics
$\Delta U = q$	Rate = $k [A]^x[B]^y[C]^z$
$w_{\text{system}} = -P\Delta V$	Rate = $k$
$\Delta E = q + w$	Rate = $k[A]$
$\Delta H = \Delta E + P\Delta V$	Rate = $k[A]^2$
$q_p = \Delta E + P\Delta V$	$[A] = -kt + [A]_0$
$\Delta E = nC_v\Delta T$	$\ln[A] = -kt + \ln[A]_0$
$\Delta H = q_p = m C_p \Delta T$	$1/[A] = kt + 1/[A]_0$
$C_p = C_v + R$	$t = [A]_0 / 2k$
$\Delta H_{\text{rxn}}^\circ = \Delta n_p \Delta H_f^\circ(\text{products}) -$ $\Delta n_r \Delta H_f^\circ(\text{reactants})$	$\ln 2 = kt$
$q_{\text{rev}} = -w_{\text{max}} = nRT \ln(V_2/V_1)$	$k = Ae^{-E_a/RT}$
$\Delta S = q_{\text{rev}} / T$	$\ln\left(\frac{k_1}{k_2}\right) = \frac{E_a}{R} \left(\frac{1}{T_2} - \frac{1}{T_1}\right)$
$\Delta S_{T_1-T_2} = nC_p \ln(T_2/T_1)$	Bonding
$\Delta S_{T_1-T_2} = nC_v \ln(T_2/T_1)$	$DE = k(Q_1Q_2 / r)$
$\Delta S_{\text{surroundings}}^\circ = \frac{q_{\text{surroundings}}}{T} = \frac{-\Delta H_{\text{sys}}}{T}$	$\Delta H_{\text{rxn}} = \sum n_p D(\text{reactants}) -$ $\sum n_r D(\text{products})$
$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$	Formal Charge = #valence $e^-$ in free atom - #lone pair $e^-$ - 1/2(# bonding $e^-$ )
$\Delta G = \Delta G^\circ + RT \ln(Q)$	
$\Delta G^\circ = -RT \ln(K)$	

### Data For Water and Other Constants

Density $d_{\text{water}} = 1.00 \text{ g/mL (25}^\circ\text{C)}$	$C_{\text{water}} = 4.184 \text{ J g}^{-1} \text{ }^\circ\text{C}^{-1}$ (liquid)
Avogadro's Number $N$	$6.022 \times 10^{23} \text{ mol}^{-1}$
Boltzmann's constant $k$	$1.30866 \times 10^{-23} \text{ J} \cdot \text{K}^{-1}$
Faraday's constant $F$	96,485 $\text{C} \cdot \text{mol}^{-1}$
Gas constant $R$	8.31451 $\text{J} \cdot \text{K}^{-1} \cdot \text{mol}^{-1}$
	0.08206 $\text{L} \cdot \text{atm} \cdot \text{K}^{-1} \cdot \text{mol}^{-1}$
Mass of Electron $m_e$	$9.10938188 \times 10^{-31} \text{ kg}$
Planck's constant $h$	$6.62608 \times 10^{-34} \text{ J} \cdot \text{s}$
Speed of Light $c$	$2.99792458 \times 10^8 \text{ m} \cdot \text{s}^{-1}$

$$K_w = 1.00 \times 10^{-14}$$

$$\text{General Information: } x = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a}$$

