

CHM 1300 and
CHM 1310 B

DATE: December 20, 2004
9:30 am

Dr. Rashmi Venkateswaran

TIME: 3 hours

Page 1 of 20

STUDENT NAME: (Print) _____

STUDENT NUMBER: _____

COURSE CODE: (Please circle) CHM 1300 CHM 1310B

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).
3. Equations and constants are given on pages 18 and 19. A periodic table of the elements is included (page 20).
4. **Use the correct number of significant figures at all times.**
5. Marks allotted to each question are shown in brackets.
7. Use the back of sheets for rough work. Three extra (blank) pages (15, 16 and 17) 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.

Quest.	1	2	3	4	5	6	7	8	9	10	
Marks	/10	/5	/20	/10	/13	/4	/8	/10	/10	/10	/100

- (10) 1. (a) The human eye can detect as little as 2.35×10^{-18} J of green light of wavelength 510 nm. Calculate the minimum number of photons that can be detected by the human eye.

$$E = h\nu = \frac{hc}{\lambda} = \frac{(6.626 \times 10^{-34} \text{ J s})(3.00 \times 10^8 \frac{\text{m}}{\text{s}})}{510 \times 10^{-9} \text{ m}}$$

$$= 3.90 \times 10^{-19} \text{ J per photon}$$

$$\begin{array}{l} 3.90 \times 10^{-19} \text{ J} \rightarrow 1 \text{ photon} \\ 2.35 \times 10^{-18} \text{ J} \rightarrow x \text{ photons} \end{array}$$

$$x = \frac{2.35 \times 10^{-18} \text{ J}}{3.90 \times 10^{-19} \text{ J/photon}} = \boxed{6.0 \text{ photons}}$$

- (b) In a photoelectric effect experiment, light of energy 6.00×10^{-19} J is absorbed by a metal, and the ejected electrons have a maximum kinetic energy of 2.70×10^{-19} J. Calculate the following:

- i) the binding energy of the electrons in the metal

$$\begin{aligned} E_{\text{kin}} &= E_{\text{hw}} - BE \\ BE &= E_{\text{hw}} - E_{\text{kin}} \\ &= 6.00 \times 10^{-19} \text{ J} - 2.70 \times 10^{-19} \text{ J} \\ &= \boxed{3.30 \times 10^{-19} \text{ J}} \end{aligned}$$

- ii) the wavelength of the light

$$\begin{aligned} E &= \frac{hc}{\lambda} & \lambda &= \frac{hc}{E} = \frac{(6.626 \times 10^{-34} \text{ J s})(3.00 \times 10^8 \frac{\text{m}}{\text{s}})}{6.00 \times 10^{-19} \text{ J}} \\ & & &= 3.31 \times 10^{-7} \text{ m} \\ & & &= \boxed{331 \text{ nm}} \end{aligned}$$

- iii) the wavelength of the electrons

$$E = \frac{1}{2} mu^2 \Rightarrow u = \sqrt{\frac{2E}{m}} = \sqrt{2(2.70 \times 10^{-19} \text{ J})} = 7.70 \times 10^5 \frac{\text{m}}{\text{s}}$$

$$\lambda = \frac{h}{mv} = \frac{6.626 \times 10^{-34} \text{ J}\cdot\text{s}}{(9.10938188 \times 10^{-31} \text{ Kg})(7.70 \times 10^5 \frac{\text{m}}{\text{s}})} = 9.45 \times 10^{-10} \text{ m} = \underline{\underline{9.45 \text{ \AA}}}$$

(e) What mass of water could have its temperature raised by 5.0°C by a mole of photons that have a wavelength of 600 nm ?

$$E = \frac{hc}{\lambda} = \frac{(6.626 \times 10^{-34} \text{ J s}) (3.00 \times 10^8 \frac{\text{m}}{\text{s}})}{600 \times 10^{-9} \text{ m}} = 3.31 \times 10^{-19} \frac{\text{J}}{\text{photon}}$$

$$\therefore E_{\text{TOT}} = \left(\frac{E}{\text{photon}} \right) (\# \text{ photons}) = \left(3.31 \times 10^{-19} \frac{\text{J}}{\text{photon}} \right) (6.022 \times 10^{23} \text{ photons})$$

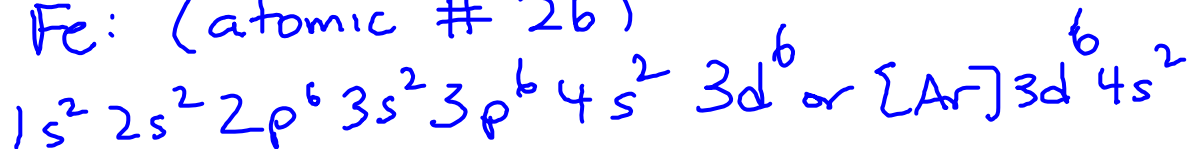
$$= 2.00 \times 10^5 \text{ J/mol}$$

$$E = mc \Delta T \Rightarrow 2.00 \times 10^5 \frac{\text{J}}{\text{mol}} = m (4.184 \frac{\text{J}}{\text{g}^\circ\text{C}}) (5.0^\circ\text{C})$$

$$m = 9.54 \times 10^3 \text{ g} \\ = \boxed{9.54 \text{ kg}}$$

- (5) 2. What are the quantum numbers of the electrons that are lost by an atom of iron when it forms the ion Fe^{2+} ? Why does it lose these particular electrons?

Fe: (atomic # 26)



when it loses $1 e^-$, it is $4s^2 3d^5$
but the second e^- is a $4s$ electron

\Rightarrow so qn's are $n=3 \quad l=2 \quad m_l = -2 \quad m_s = -\frac{1}{2}$
 $n=4 \quad l=0 \quad m_l = 0 \quad m_s = -\frac{1}{2}$

It loses these particular electrons so it forms a stable half filled orbital for both $4s$ and $3d$.

(20) 3. A certain element X was found to form three compounds with chlorine having the formulae XCl_2 , XCl_4 , and XCl_6 . One of its oxides has the formula XO_3 , and X reacts with sodium to form the compound Na_2X .

(a) Is X a metal or a non metal?

non metal

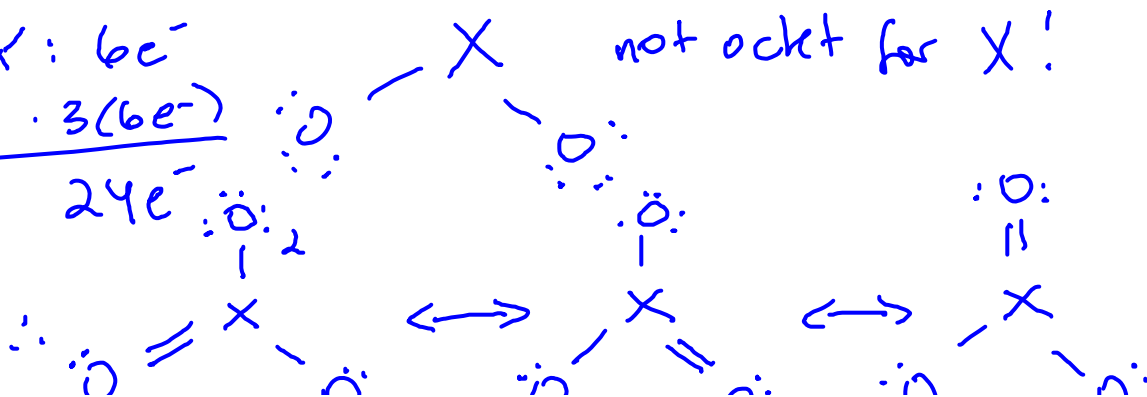
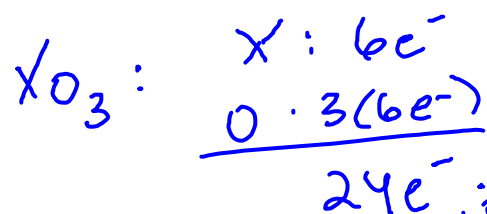
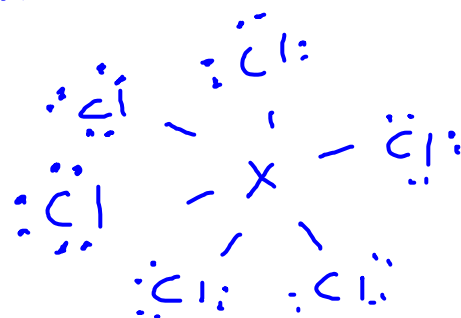
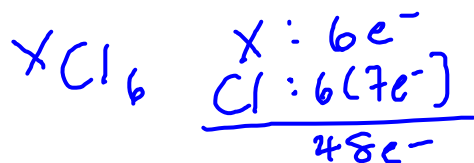
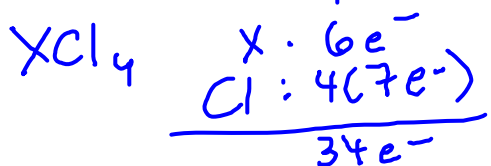
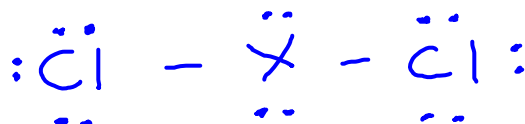
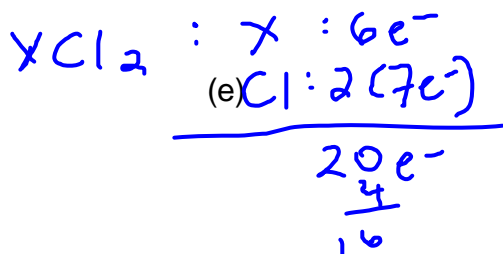
(b) In which group in the periodic table is X located?

Group 6A or 16

(c) In which rows (periods) in the periodic table could X possibly be located?

2, 3, 4, 5 or 6

(d) Draw Lewis structures for XCl_2 , XCl_4 , and XCl_6 and XO_3 . Where possible, follow the octet rule. Which of the structures has (have) multiple bonding? Are there any resonance structures to consider?



What do we expect the shapes (electron geometries) of the 4 molecules drawn above to be? Which molecules would be polar? (simply state the names of the shapes...you may draw them only if you wish).

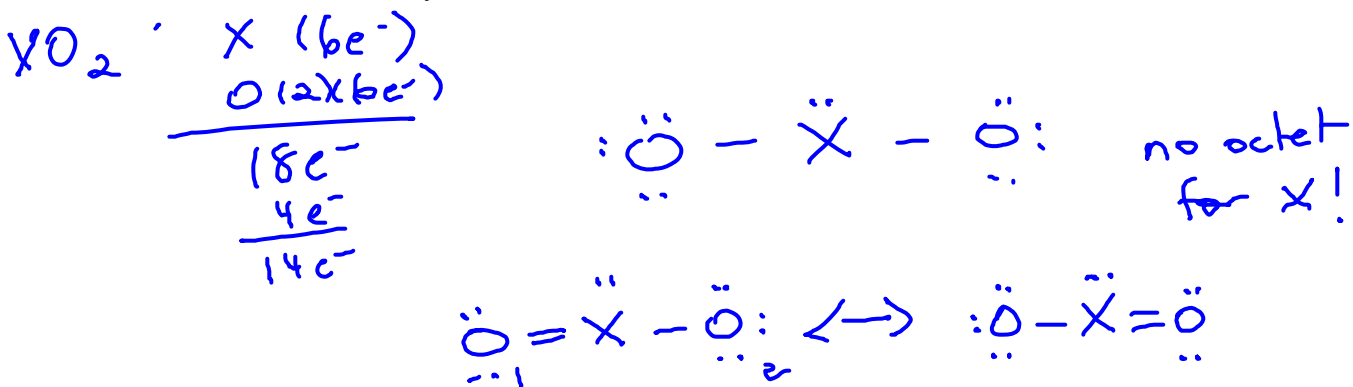
XCl_2 : MG: tetrahedral shape: bent

XCl_3 : MG: trigonal bipyramidal shape: see saw

XCl_6 : MG and shape: octahedral

XO_3 : MG and shape: trigonal planar

- (f) The element X also forms the oxide XO_2 . Draw the Lewis structure for this molecule that obeys the octet rule.



- (g) Assign formal charges to the atoms in the Lewis structures for XO_2 and XO_3 .

XO_2 :

$$\begin{aligned} \text{FC}_X &= 6 - 2 = 4 \\ \text{FC}_{\text{O}_1} &= 6 - 4 = 2 \\ \text{FC}_{\text{O}_2} &= 6 - 6 - 1 = -1 \end{aligned}$$

XO_3

$$\begin{aligned} \text{FC}_X &= 6 - 0 - 4 = 2 \\ \text{FC}_{\text{O}_1} &= 6 - 4 - 2 = 0 \\ \text{FC}_{\text{O}_2} = \text{FC}_{\text{O}_3} &= 6 - 6 - 1 = -1 \end{aligned}$$

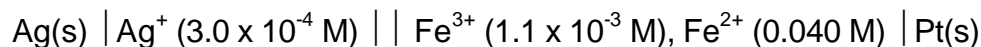
- (h) If X were to form a compound with Aluminum, what would be its formula?



- (i) If X were in Period 5, what would be the electron configuration of X expressed using a noble gas core configuration?



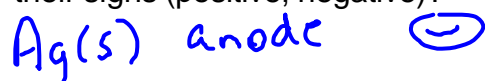
- (10) 4. (a) Consider the following galvanic cell:



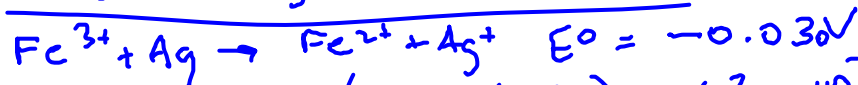
- i) Write the balanced equation for the spontaneous cell reaction.



- ii) Which electrode is the anode and which is the cathode and what are their signs (positive, negative)?



- iii) Calculate the cell potential given the following standard potentials:



$$E = E^\circ - \frac{RT}{nF} \ln Q = -0.030 \text{ V} - \frac{(8.314 \text{ J K}^{-1} \text{ mol}^{-1})(298)}{(1)(96485)} \ln \left(\frac{(3.0 \times 10^{-4})(0.040)}{(1.1 \times 10^{-3})} \right)$$

$$= \boxed{0.086 \text{ V}}$$

- (b) A solution containing vanadium in an unknown oxidation state was electrolyzed with a current of 1.50 A for 30.0 min. It was found that 0.475 g of V was deposited on the cathode. What was the original oxidation state of the vanadium ion?

$$Q = It = (1.50 \text{ A})(30.0 \text{ min}) \left(\frac{60 \text{ s}}{\text{min}} \right) = 2.70 \times 10^3 \text{ C}$$



$$\times = 2.70 \times 10^3 / 96485 = 2.80 \times 10^{-2} \text{ mole}^-$$

$$n_V = \frac{m}{MM} = \frac{0.475 \text{ g}}{50.9415 \frac{\text{g}}{\text{mol}}} = 9.32 \times 10^{-3} \text{ mol}$$



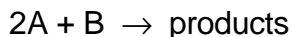
$$n = \frac{2.80 \times 10^{-2} \text{ mol}}{9.32 \times 10^{-3} \text{ mol}} = 3$$

$$\therefore x \text{ mole } \text{e}^- \rightarrow 1 \text{ mol V}$$

$$2.80 \times 10^{-2} \text{ mole}^- \rightarrow 9.32 \times 10^{-3} \text{ mol}$$

oxidation state is $(+3)$

(13) 5. (a) A reaction that has the stoichiometry below was found to yield the following data:



The initial rate of this reaction was determined for several concentrations of HgCl_2 and $\text{C}_2\text{O}_4^{2-}$ and the following rate data were obtained for the rate of disappearance of $\text{C}_2\text{O}_4^{2-}$:

Experiment	[A](M)	[B](M)	Initial Rate (M/s)
1	0.020	0.030	0.0150
2	0.025	0.030	0.0188
3	0.025	0.040	0.0334

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

$$\left(\frac{0.025}{0.020}\right)^x = \left(\frac{0.0188}{0.0150}\right) \quad \left(\frac{0.040}{0.030}\right)^y = \left(\frac{0.0334}{0.0188}\right)$$

$$1.25^x = 1.25 \quad 1.33^y = 1.78$$

$$x = 1 \quad y = 2$$

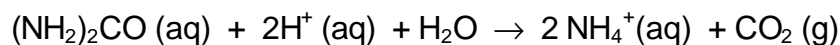
$$\therefore \text{Rate} = k[A][B]^2$$

(ii) What is the rate constant for this reaction with the correct units?

$$0.0150 \frac{\text{M}}{\text{s}} = k (0.020 \text{ M})(0.030 \text{ M})^2$$

$$k = 8.33 \times 10^{-2} \text{ M}^{-2} \text{ s}^{-1}$$

(b) The decomposition of urea, $(\text{NH}_2)_2\text{CO}$, in 0.10 M HCl follows the equation:



At 60.0 °C, $k = 5.84 \times 10^{-6} \text{ min}^{-1}$ and at 70.0 °C, we find that $k = 2.25 \times 10^{-5} \text{ min}^{-1}$. If this reaction is run at 80.0 °C starting with a urea concentration of 0.0020 M, how many minutes will it take for the urea concentration to drop to 0.0012 M?

$$\ln \frac{k_1}{k_2} = \frac{E_a}{R} \left(\frac{1}{T_2} - \frac{1}{T_1} \right)$$

$$\ln \left(\frac{5.84 \times 10^{-6}}{2.25 \times 10^{-5}} \right) = \frac{E_a}{8.314} \left(\frac{1}{70.0 + 273} - \frac{1}{60.0 + 273} \right)$$

$$\ln \frac{[A_t]}{[A_0]} = -kt$$

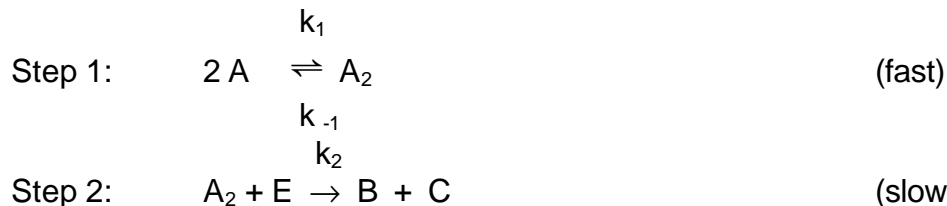
$$E_a = 1.28 \times 10^5 \text{ J}$$

$$\ln \left(\frac{0.3812}{0.8820} \right) = - \frac{(1.28 \times 10^5 \text{ J})}{8.314 \text{ J mol}^{-1} \text{ K}^{-1}} \left(\frac{1}{343} - \frac{1}{353} \right)$$

$$t = 6.36 \times 10^3 \text{ s} = \boxed{106 \text{ min}}$$

$$k_{350} = 8.03 \times 10^{-5}$$

(c) Suppose a reaction occurs with the mechanism:



(i) What is the overall reaction?



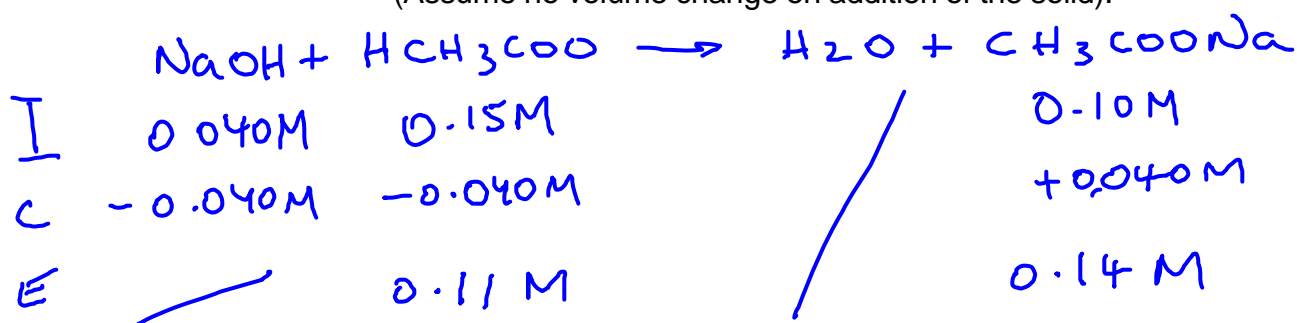
(ii) Using the definition of equilibrium, determine the rate expression for the overall reaction in terms of the reactants only.

$$k_1 [A]^2 = k_{-1} [A_2]$$

$$\text{Rate} = k_2 [A_2] [E] = k_2 \frac{k_1 [A]^2 [E]}{k_{-1}}$$

$$= k [A]^2 [E]$$

If 0.020 mol of NaOH were added to 500.0 mL of a sodium acetate-acetic acid buffer that contains 0.10 M sodium acetate and 0.15 M acetic acid, by how many pH units will the pH of the buffer change? (Assume no volume change on addition of the solid).



$$\text{Original pH} = \text{pK} + \log \frac{\text{A}^-}{\text{HA}} = -\log(1.8 \times 10^{-5}) + \log \frac{0.10}{0.15} = 4.57$$

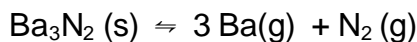
$$\text{New pH} = \text{pK} + \log \frac{\text{A}^-}{\text{HA}} = -\log(1.8 \times 10^{-5}) + \log \frac{0.14}{0.11} = 4.85$$

$$\therefore \text{diff in pH} = 4.85 - 4.57 = \boxed{0.28 \text{ pH units}}$$

(c) For the reaction $\text{NO}_2(\text{g}) + \text{SO}_2(\text{g}) \rightleftharpoons \text{NO}(\text{g}) + \text{SO}_3(\text{g})$, $K_c = 3.6$ and $\Delta H^\circ = -41.8 \text{ kJ/mol}$. Answer, in 1-3 words, how the equilibrium concentration of NO will be affected if:

- i) More NO_2 is added to the container. increase
- ii) Some SO_3 is removed from the container. increase
- iii) The temperature of the reaction mixture is raised. decrease
- iv) Some SO_2 is removed from the mixture. decrease
- v) The pressure of the gas mixture is lowered by expanding the volume to 10.0 L. no change

- (10) 8. (a) Barium nitride vaporizes slightly at high temperature by means of the dissociation:



At 1000. K, the equilibrium constant is 4.5×10^{-19} . At 1200. K, the equilibrium constant is 6.2×10^{-12} .

- i) Determine the value of ΔH° for the reaction.

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

$$\ln \left(\frac{4.5 \times 10^{-19}}{6.2 \times 10^{-12}} \right) = \frac{\Delta H^\circ}{8314} \left(\frac{1}{1200} - \frac{1}{1000} \right)$$

$$\Delta H^\circ = 8.20 \times 10^5 \text{ J/mol}$$

- ii) What approximation was made in part (i)?

We assume ΔH° is constant in this temperature range.

- (b) Suppose 61.0 g of hot metal that is initially at 120.0 °C is plunged into 100.0 g of water that is initially at 20.00 °C. The final temperature of the mixture is 26.39 °C and no heat is lost to the surroundings. Calculate the specific heat capacity of the metal, given that the specific heat capacity of water is $4.184 \text{ J } ^\circ\text{C}^{-1}\text{g}^{-1}$.

$$-q_{\text{metal}} = q_{\text{water}}$$

$$-mc \Delta T_{\text{met}} = mc \Delta T_w$$

$$-(61.0 \text{ g})(c)(26.39 - 120.0)^\circ\text{C} = (100.0 \text{ g}) \left(4.184 \frac{\text{J}}{\text{g}^\circ\text{C}} \right) (26.39 - 20.00)^\circ\text{C}$$

$$c = 0.468 \frac{\text{J}}{\text{g}^\circ\text{C}} \quad \text{for the unknown metal.}$$

- (10) 9. A sample of a new antimalarial drug with a mass of 0.2394 g was made to undergo a series of reactions that changed all of the nitrogen in the drug into N_2 . The nitrogen gas had a volume of 18.90 mL when collected over water at 23.80 °C and a pressure of 746.0 Torr. At 23.80 °C, the vapour pressure of water is 22.110 Torr.

(a) Calculate the percentage of nitrogen in the sample.

$$P_{N_2} = 746.0 - 22.110 = 723.9 \text{ Torr}$$

$$n_{N_2} = \frac{PV}{RT} = \frac{\left(\frac{723.9}{760}\right)(0.01890 \text{ L})}{\left(0.08206 \frac{\text{Latm}}{\text{molK}}\right)(23.80 + 273 \text{ K})} = 7.39 \times 10^{-4} \text{ mol}$$

$$\text{mol N} = 2 \text{ mol } N_2 = 1.48 \times 10^{-3} \text{ mol}$$

$$\text{mass N} = n \text{ MM} = (1.48 \times 10^{-3})(14) = 2.07 \times 10^{-2} \text{ g}$$

$$\text{mass } \% \text{ N} = \frac{2.07 \times 10^{-2} \text{ g}}{0.2394 \text{ g}} = \boxed{8.64 \%}$$

$$\text{mass of N in } 6.478 \text{ mg sample} = 5.60 \times 10^{-4} \text{ g}$$

$$n_N = \frac{5.60 \times 10^{-4} \text{ g}}{14} = 4 \times 10^{-5}$$

(b) When 6.478 mg of the compound was burned in pure oxygen, 17.57 mg of CO_2 and 4.319 mg of H_2O were obtained. What are the percentages of C and H in this compound?

$$n_{CO_2} = \frac{17.57 \times 10^{-3} \text{ g}}{44 \text{ g/mol}} = 3.99 \times 10^{-4} \text{ mol}$$

$$n_C = n_{CO_2} = 3.99 \times 10^{-4} \text{ mol} \Rightarrow M_C = n_C \text{ MM}_C = (3.99 \times 10^{-4} \text{ mol})(12 \frac{\text{g}}{\text{mol}}) = 4.79 \times 10^{-3} \text{ g}$$

$$\text{mass } \% \text{ C} = \frac{4.79 \times 10^{-3}}{6.478 \times 10^{-3}} \times 100\% = 73.97\%$$

$$n_{H_2O} = \frac{4.319 \times 10^{-3} \text{ g}}{18 \text{ g/mol}} = 2.399 \times 10^{-4} \text{ mol}$$

$$n_H = 2n_{H_2O} = 4.799 \times 10^{-4} \text{ mol} \Rightarrow M_H = n_H \text{ MM}_H = 4.799 \times 10^{-4} \text{ g}$$

$$\text{mass } \% \text{ H} = \frac{4.799 \times 10^{-4} \text{ g}}{6.478 \times 10^{-3} \text{ g}} \times 100\% = 7.41\%$$

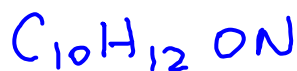
- (c) Assuming that any undetermined element is oxygen, write an empirical formula for the compound.

$$\text{mass \% O} = 100 - 73.97 - 7.41 - 8.64 = 9.98\%$$

$$\therefore \text{in } 6.778 \text{ mg sample, } m_{\text{O}} = 6.465 \times 10^{-4} \text{ g}$$

$$n_{\text{O}} = \frac{6.465 \times 10^{-4} \text{ g}}{16 \text{ g/mol}} = 4.04 \times 10^{-5} \text{ mol}$$

$$\begin{array}{cccc} \text{C} & \text{H} & \text{O} & \text{N} \\ \frac{3.99 \times 10^{-4}}{4.0 \times 10^{-5}} & \frac{4.799 \times 10^{-4}}{4.0 \times 10^{-5}} & \frac{4.04 \times 10^{-5}}{4.0 \times 10^{-5}} & \frac{4.0 \times 10^{-5}}{4.0 \times 10^{-5}} \end{array}$$



- (d) The molecular mass of the compound was found to be 324 g/mol. What is its molecular formula?

$$\begin{aligned} \text{MM}_{\text{emp form}} &= 10(12) + 12(1) + 1(16) + 1(14) \\ &= 162 \end{aligned}$$

$$\text{MM}_{\text{mol form}} = 324$$

$$\therefore \text{ratio} = 2$$

$$\therefore \text{Molecular formula} = \text{C}_{20}\text{H}_{24}\text{O}_2\text{N}_2$$

(10) 10. Choose one of the following topics: quantum chemistry, corrosion, batteries, acids and bases, equilibrium, kinetics, gas laws, or thermodynamics. In a letter of minimum length 1 page and using at least 3 chemical concepts (clearly explained), write to your mother and explain why the particular topic you chose will have an impact on your personal and professional life.

no specific answer

Equations and Constants

<p style="text-align: center;"><u>Gas Law</u></p> $PV = nRT$ $P_{\text{Total}} = P_1 + P_2 + P_3 + \dots$ $d = m/V = P(\text{MM}) / RT$ $KE = (1/2)mv_{\text{av}}^2$ $\sqrt{v^2} = \sqrt{\frac{3RT}{M}}$ $\frac{\text{Rate}_A}{\text{Rate}_B} = \sqrt{\frac{M_b}{M_a}}$ $P + \frac{n^2 a}{V^2}(V - nb) = nRT$ <p>1 atm = 760 Torr = 760 mm Hg = 1.01325 x 10⁵ Pa</p> <p style="text-align: center;"><u>Equilibrium</u></p> $K_p = K(RT)^{\Delta n}$ $K = [C]^c [D]^d / [A]^a [B]^b$ $K_{\text{sp}} = [C]^c [D]^d$ <p style="text-align: center;"><u>Acid/Base</u></p> $\text{pOH} = -\log [\text{OH}^-]$ $\text{pH} = -\log [\text{H}^+]$ $\text{pH} + \text{pOH} = 14$ $K_a \times K_b = K_w$ $\text{pH} = \text{p}K_a + \log \left[\frac{[\text{A}^-]}{[\text{HA}]}\right]$ $\text{pH} = \frac{\text{p}K_{a1} + \text{p}K_{a2}}{2}$ <p>Concentration (M) = n/V</p>	<p style="text-align: center;"><u>Electrochemistry</u></p> $\Delta G^\circ = -nFE^\circ$ $E = E^\circ - \frac{RT}{nF} \ln(Q)$ <p>Or $E = E^\circ - \frac{0.059}{n} \log(Q)$ at 25°C</p> $Q = I \cdot t$ <p style="text-align: center;"><u>Quantum Mechanics</u></p> $? \cdot ? = c$ $E = h \cdot ?$ $m = \frac{h}{\lambda c}$ <p>Energy of state = -2.178x10⁻¹⁸ J / n²</p> $\Delta x \cdot \Delta p \geq h / 4\pi$ $E = -C(1/n^2)$ <p style="text-align: center;"><u>Liquids and Colligative Properties</u></p> $\ln(P_1/P_2) = \Delta H^\circ / R (1/T_2 - 1/T_1)$ $P_{\text{solution}} = X_{\text{solvent}} \cdot P_{\text{solvent}}^\circ$ $\Delta T_{\text{BP}} = K_{\text{BP}} \cdot m$ $\Delta T_{\text{FP}} = K_{\text{FP}} \cdot m$ $\Delta T = K \cdot m \cdot i$ $? = cRT$ <p>molality = mol solute/mass solvent (kg)</p>
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Data for Question 9	ΔG° (kJ / mol) at 25 °C	
SO ₂ (g)	-300.4	
H ₂ S (g)	-33.01	
H ₂ O (g)	-228.57	

<u>Thermodynamics</u>	<u>Kinetics</u>
$\Delta U = q$	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 = $k[A]^2$
$q_p = \Delta E + P \Delta V$	$[A] = -kt + [A]_0$
$\Delta E = nC_v \Delta T$	$\ln[A] = -kt + \ln[A]_0$
$\Delta H = q_p = m C_p \Delta T$	$1/[A] = kt + 1/[A]_0$
$C_p = C_v + R$	$t = [A]_0 / 2k$
$\Delta H_{\text{rxn}}^\circ = \Delta n_p \Delta H_f^\circ(\text{products})?$	$\ln 2 = kt$
$\Delta n_r \Delta H_f^\circ(\text{reactants})$	$k = A e^{-E_a/RT}$
$q_{\text{rev}} = -w_{\text{max}} = nRT \ln(V_2/V_1)$	$\ln(k_1/k_2) = E_a/R (1/T_2 - 1/T_1)$
$\Delta S = q_{\text{rev}} / T$	<u>Bonding</u>
$\Delta S_{T_1-T_2} = nC_p \ln(T_2/T_1)$	$DE = k (Q_1 Q_2 / r)$
$\Delta S_{T_1-T_2} = nC_v \ln(T_2/T_1)$	$\Delta H_{\text{rxn}} = \sum n_p D(\text{reactants}) -$
$\Delta S_{\text{surroundings}}^\circ = \frac{q_{\text{surroundings}}}{T} = \frac{-\Delta H_{\text{sys}}}{T}$	$\sum n_r D(\text{products})$
$\Delta G^\circ = \Delta H^\circ - T \Delta S^\circ$	Formal Charge = #valence e^- in free atom - #lone pair e^- - 1/2(# bonding e^-)
$\Delta G = \Delta G^\circ + RT \ln(Q)$	
$\Delta G^\circ = -RT \ln(K)$	
$\ln(K_1/K_2) = \Delta H^\circ/R (1/T_2 - 1/T_1)$	

Data For Water and Other Constants

Density $d = 1.00 \text{ g/mL (25}^\circ\text{C)}$	$C_p = 2.01 \text{ J g}^{-1} \text{ }^\circ\text{C}^{-1} \text{ (gas)}$
$C = 2.13 \text{ J g}^{-1} \text{ }^\circ\text{C}^{-1} \text{ (solid)}$	$\Delta H_{\text{vap}}^\circ = 40.7 \text{ kJ mol}^{-1}$
$C = 4.18 \text{ J g}^{-1} \text{ }^\circ\text{C}^{-1} \text{ (liquid)}$	$\Delta H_{\text{fus}}^\circ = 6.02 \text{ kJ mol}^{-1}$
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	$8.31451 \text{ J} \cdot \text{K}^{-1} \cdot \text{mol}^{-1}$
	$0.08206 \text{ L} \cdot \text{atm} \cdot \text{K}^{-1} \cdot \text{mol}^{-1}$
Planck's constant h	$6.62608 \times 10^{-34} \text{ J} \cdot \text{s}$
Speed of Light c	$2.99792458 \times 10^8 \text{ m} \cdot \text{s}^{-1}$

