

CHEM 1110

ANSWERS TO STOICHIOMETRY PROBLEM SET

$$1. \quad \text{moles of SCl}_2 = 4.00 \text{ g SCl}_2 \times \frac{1 \text{ mol SCl}_2}{103.1 \text{ g SCl}_2} = 0.03880 \text{ mol SCl}_2$$

$$\text{moles NaF} = 2.00 \text{ g NaF} \times \frac{1 \text{ mol NaF}}{42.0 \text{ g NaF}} = 0.04762 \text{ mole NaF}$$

moles SCl₂ needed to react with 0.04762 mol NaF

$$= 0.04762 \text{ mol NaF} \times \frac{3 \text{ mol SCl}_2}{4 \text{ mol NaF}} = 0.03572 \text{ mol SCl}_2$$

BUT, 0.03880 mol SCl₂ is available; hence SCl₂ is in EXCESS and NaF is LIMITING.

$$\text{moles SF}_4 = 0.04762 \text{ mol NaF} \times \frac{1 \text{ mol SF}_4}{4 \text{ mol NaF}} = 0.01191 \text{ mol SF}_4$$

$$\text{mass SF}_4 = 0.01191 \text{ mol SF}_4 \times \frac{108.1 \text{ g SF}_4}{1 \text{ mol SF}_4} = \mathbf{1.29 \text{ g SF}_4}$$

$$2. \quad \text{moles BF}_3 = 54.0 \text{ g BF}_3 \times \frac{1 \text{ mol BF}_3}{67.8 \text{ g BF}_3} = 0.7965 \text{ mol BF}_3$$

$$\text{mass NaBH}_4 = 25.0 \text{ g "85.0\% NaBH}_4" \times \frac{85.0 \text{ g NaBH}_4}{100 \text{ g "85.0\% NaBH}_4"} = 21.25 \text{ g NaBH}_4$$

$$\text{moles NaBH}_4 = 21.25 \text{ g NaBH}_4 \times \frac{1 \text{ mol NaBH}_4}{37.8 \text{ g NaBH}_4} = 0.5622 \text{ mol NaBH}_4$$

moles NaBH₄ needed to react with with 0.7965 mol BF₃

$$= 0.7965 \text{ mol BF}_3 \times \frac{3 \text{ mol NaBH}_4}{4 \text{ mol BF}_3} = 0.5974 \text{ mol NaBH}_4$$

BUT, only 0.5622 mol NaBH₄ is available; hence NaBH₄ is the LIMITING REACTANT.

$$\text{moles B}_2\text{H}_6 = 0.5622 \text{ mol NaBH}_4 \times \frac{2 \text{ mol B}_2\text{H}_6}{3 \text{ mol NaBH}_4} = 0.3748 \text{ mol B}_2\text{H}_6$$

$$\text{mass B}_2\text{H}_6 = 0.3748 \text{ mol B}_2\text{H}_6 \times \frac{27.7 \text{ g B}_2\text{H}_6}{1 \text{ mol B}_2\text{H}_6} = \mathbf{10.4 \text{ g B}_2\text{H}_6}$$

$$3. \quad \text{moles O}_2 = 0.655 \text{ L O}_2 \times \frac{1 \text{ mol O}_2}{22.4 \text{ L O}_2} = 0.02924 \text{ mol O}_2$$

$$\text{moles KO}_2 = 0.02924 \text{ mol O}_2 \times \frac{4 \text{ mol KO}_2}{3 \text{ mol O}_2} = 0.03899 \text{ mole KO}_2$$

$$\text{mass KO}_2 = 0.03899 \text{ mol KO}_2 \times \frac{71.1 \text{ g KO}_2}{1 \text{ mol KO}_2} = 2.772 \text{ g KO}_2$$

$$\% \text{ purity} = \frac{2.772}{3.30} \times 100 = \mathbf{84.0 \%}$$

4. MW of C₁₇H₂₁O₄N (cocaine; "coc") = 303.2 g/mole
 MW of C₁₂H₂₂O₁₁ (sucrose; "suc") = 342.2 g/mole

1 mole of C₁₇H₂₁O₄N produces 17 moles of CO₂ on combustion

1 mole of C₁₂H₂₂O₁₁ produces 12 moles of CO₂ on combustion

Let x = mass (mg) of cocaine; $1-x$ = mass (mg) of sucrose

mass of CO₂ produced from cocaine =

$$\frac{x}{303.2} \text{ mmol coc} \times \frac{17 \text{ mmol CO}_2}{1 \text{ mmol coc}} \times \frac{44.01 \text{ mg CO}_2}{1 \text{ mmol CO}_2} = 2.468x \text{ mg CO}_2$$

mass of CO₂ produced from sucrose =

$$\frac{(1-x)}{342.2} \text{ mmol suc} \times \frac{12 \text{ mmol CO}_2}{1 \text{ mmol suc}} \times \frac{44.01 \text{ mg CO}_2}{1 \text{ mmol CO}_2} = 1.543 - 1.543x$$

$$\text{total mass of CO}_2 = 2.468x + 1.543 - 1.543x = 1.75$$

$$(2.468 - 1.543)x = 1.75 - 1.543$$

$$0.928x = 0.207$$

$$x = 0.22 \text{ mg}$$

$$\% \text{ cocaine} = \frac{0.22 \text{ mg cocaine}}{1.00 \text{ mg mixture}} \times 100 = \mathbf{22\%}$$

5. 86 proof is equivalent to 43 mL of ethanol (EtOH) in 100 mL of solution.
 Consider 100 mL of solution \equiv 0.100 L

$$\text{moles EtOH} = 43 \text{ mL EtOH} \times \frac{0.79 \text{ g EtOH}}{1 \text{ mL EtOH}} \times \frac{1 \text{ mol EtOH}}{46.0 \text{ g EtOH}} = 0.738 \text{ mol}$$

$$\text{molarity} = \frac{0.738 \text{ mol EtOH}}{0.100 \text{ L}} = \mathbf{7.4 M}$$

6. Since I_2 is in excess, SO_3^{2-} is limiting.
 No. of moles SO_3^{2-} oxidized = No. of moles $Na_2SO_3 \cdot XH_2O$ originally present

$$= 0.0400 \text{ L NaOH sol.} \times \frac{0.100 \text{ mol NaOH}}{1 \text{ L NaOH sol.}} \times \frac{1 \text{ mol OH}^-}{1 \text{ mol NaOH}} \times \frac{1 \text{ mol H}^+}{1 \text{ mol OH}^-}$$

$$\times \frac{1 \text{ mol } SO_3^{2-}}{2 \text{ mol H}^+} \times \frac{1 \text{ mol } SO_3^{2-}}{1 \text{ mol } Na_2SO_3 \cdot XH_2O} = 2.00 \times 10^{-3} \text{ mol } Na_2SO_3 \cdot XH_2O$$

$$\text{Molar mass of } Na_2SO_3 \cdot XH_2O = \frac{0.4322 \text{ g } Na_2SO_3 \cdot XH_2O}{2.00 \times 10^{-3} \text{ mol } Na_2SO_3 \cdot XH_2O} = 216 \text{ g/mol}$$

$$X = \frac{\text{Molar mass } Na_2SO_3 \cdot XH_2O - \text{Molar mass } Na_2SO_3}{\text{Molar mass } H_2O}$$

$$= \frac{216 - 126}{18.0} = 5$$

Therefore, formula is **$Na_2SO_3 \cdot 5H_2O$**

7. Let X g = mass of Cu in alloy and Y g = mass of Ag in alloy

mass of NO_2 formed from X g Cu =

$$X \text{ g Cu} \times \frac{1 \text{ mol Cu}}{63.55 \text{ g Cu}} \times \frac{2 \text{ mol } NO_2}{1 \text{ mol Cu}} \times \frac{46.01 \text{ g } NO_2}{1 \text{ mol } NO_2} = 1.448 X$$

mass of NO_2 formed from Y g Ag =

$$Y \text{ g Ag} \times \frac{1 \text{ mol Ag}}{107.9 \text{ g Ag}} \times \frac{1 \text{ mol } NO_2}{1 \text{ mol Ag}} \times \frac{46.01 \text{ g } NO_2}{1 \text{ mol } NO_2} = 0.4264 Y$$

$$\text{total mass of } NO_2 \text{ formed} = 1.448 X + 0.4264 Y = 10.00 \text{ g} \quad \dots(1)$$

$$\text{But, } X + Y = 10.0 \text{ g;} \quad \text{therefore, } X = 10.0 - Y \quad \dots(2)$$

substitution of Eqn. 2 into Eqn.1 gives:

$$1.448 (10.0 - Y) + 0.4264 Y = 10.00$$

hence, Y = 4.385 g = mass of Ag

$$\% \text{ Ag by mass} = \frac{4.385 \text{ g Ag}}{10.0 \text{ g alloy}} \times 100 = \mathbf{43.85\%}$$