1. Vanillin, the dominant flavoring in vanilla, contains C, H, and O. When 1.05 g of thus substance is completely combusted, 2.43 g of CO₂ and 0.50 g of H₂O are produced. What is the empirical formula of vanillin?

Yet another combustion analysis problem!

$$0.50 \ g \ H_2O * \frac{1 \ mol \ H_2O}{18.02 \ g \ H_2O} * \frac{2 \ mol \ H}{1 \ mol \ H_2O} = 0.05549 \ mol \ H * \frac{1.008 \ g \ H}{1 \ mol \ H} = 0.0559 \ g \ H$$

$$2.43 \ g \ CO_2 * \frac{1 \ mol \ CO_2}{44.01 \ g \ CO_2} * \frac{1 \ mol \ C}{1 \ mol \ CO_2} = 0.05521 \ mol \ C * \frac{12.011 \ g \ C}{1 \ mol \ C} = 0.6632 \ g \ C$$

$$Mass\ O = Mass\ sample - Mass\ C - Mass\ H = 1.05\ g - 0.6632\ g - 0.0559\ g = 0.3309\ g\ O$$

$$0.3309 \ g \ O * \frac{1 \ mol \ O}{15.9994 \ g \ O} = 0.0207 \ mol \ O$$

Take mole ratios to find empirical formula!

$$C: \frac{0.05521}{0.0207} = 2.67$$

$$H: \frac{0.05549}{0.0207} = 2.67$$

$$0: \frac{0.0207}{0.0207} = 1$$

2.67 is equal to 8/3. Thus, multiply everything by 3 to get the empirical formula: C₈H₈O₃

2. A mixture of N₂(g) and H₂(g) reacts in a closed container to form ammonia, NH₃(g). The reaction ceases before either reactant has been totally consumed. At this stage, 3.0 mol N₂, 3.0 mol H₂, and 3.0 mol NH₃ are present. How many moles of N₂ and H₂ were present originally?

First write the balanced equation!

$$N_2(g) + 3H_2(g) \rightarrow 2NH_3(g)$$

Since we assume that we started off with zero product (zero ammonia), let's use that information to find out how much reactant was needed to generate that much product.

$$3.0 \ mol \ NH_3 * \frac{1 \ mol \ N_2}{2 \ mol \ NH_3} = 1.5 \ mol \ N_2 \ needed$$

$$3.0 \ mol \ NH_3 * \frac{3 \ mol \ H_2}{2 \ mol \ NH_3} = 4.5 \ mol \ H_2 \ needed$$

Thus, 1.5 mol N_2 and 4.5 mol H_2 were needed to bring the reaction to its current state, with 3 mol NH_3 . However, there are also current mole amounts of each of the reactants, as noted in the problem. Thus,

$$Original \ N_2 amount = 1.5 \ mol + 3.0 \ mol = 4.5 \ mol \ N_2$$

Original
$$H_2$$
amount = 4.5 mol + 3.0 mol = 7.5 mol H_2

3. A mixture containing KClO₃, K₂CO₃, KHCO₃, and KCl was heated, producing CO₂, O₂, and H₂O gases according to the following equations:

$$2KClO_3(s) \rightarrow 2KCl(s) + 3O_2(g)$$
 $2KHCO_3(s) \rightarrow K_2O(s) + H_2O(g) + 2CO_2(g)$ $K_2CO_3(s) \rightarrow K_2O(s) + CO_2(g)$

The KCl does not react under the conditions of the reaction. If 100.0 g of the mixture produces 1.80 g of H₂O, 13.20 g of CO₂, and 4.00 g of O₂, what was the composition of the original mixture? (Assume complete decomposition of the mixture.)

This is similar to the combustion analysis in some ways. Recognize that all of the O_2 produced is due to the first reaction, and all of the H_2O produced comes from the second reaction.

$$4.00\ g\ O_2*\frac{1\ mol\ O_2}{31.9988\ g\ O_2}*\frac{2\ mol\ KClO_3}{3\ mol\ O_2}*\frac{122.55\ g\ KClO_3}{1\ mol\ KClO_3}=10.21\ g\ KClO_3$$

$$1.80 \; g \; H_2O * \frac{1 \; mol \; H_2O}{18.02 \; g \; H_2O} * \frac{2 \; mol \; KHCO_3}{1 \; mol \; H_2O} * \frac{100.12 \; g \; KHCO_3}{1 \; mol \; KHCO_3} = 20.00 \; g \; KHCO_3$$

Next, we know how much CO_2 is produced, but that is produced in both the 2^{nd} and 3^{rd} reactions. So, since we just determined how much $KHCO_3$ was present, determine how much CO_2 that would have produced.

$$1.80 \ g \ H_2O * \frac{1 \ mol \ H_2O}{18.02 \ g \ H_2O} * \frac{2 \ mol \ CO_2}{1 \ mol \ H_2O} * \frac{44.01 \ g \ CO_2}{1 \ mol \ CO_2} = 8.79 \ g \ CO_2$$

Thus, the CO₂ produced from reaction 3 is:

$$13.20 g CO_2 - 8.79 g CO_2 = 4.41 g CO_2$$

Then, this can be used to find how much K₂CO₃ was present in the original mixture.

$$4.41 \ g \ CO_2 * \frac{1 \ mol \ CO_2}{44.01 \ g \ CO_2} * \frac{1 \ mol \ K_2CO_3}{1 \ mol \ CO_2} * \frac{138.21 \ g \ K_2CO_3}{1 \ mol \ K_2CO_3} = 13.84 \ g \ K_2CO_3$$

$$Mass\ KCl = Mass\ sample - Mass\ KClO_3 - Mass\ KHCO_3 - Mass\ K_2CO_3 = 100.0 - 10.21 - 20.00 - 13.84 = 56.0\ g\ KCl$$

Composition:

$$\% KCl = \frac{56.0 g}{100.0 g} = 56.0\% KCl$$

$$\% KClO_3 = \frac{10.21 g}{100.0 g} = 10.2\% KClO_3$$

$$\% KHCO_3 = \frac{20.00 g}{100.0 g} = 20.0\% KClO_3$$

$$\% K_2CO_3 = \frac{13.84 g}{100.0 g} = 13.8\% K_2CO_3$$

- 4. Give the oxidation number of each element in the following compounds:
 - a. BrO₃: O: -2; Br: +5
 - **b.** H₂SO₄: O: -2; S: +6; H: +1
 - c. CrO₄²⁻: O: -2; Cr: +6
 - d. LiAlH₄: Li: +1; Al: +3; H: -1
- 5. Your friend has heard that she can make ethanol by reacting C_2H_4 with H_2O under acidic conditions, but she's not sure how much of each starting material she needs. So she randomly mixes 101.7 g of C_2H_4 with 55.19 g of H_2O .

$$C_2H_4 + H_2O \rightarrow C_2H_6O$$

a. What is the theoretical yield of ethanol in mL (ethanol density = 0.789 g/mL)?

$$101.7 \ g \ C_2 H_4 * \frac{1 \ mol \ C_2 H_4}{28.05 \ g \ C_2 H_4} * \frac{1 \ mol \ C_2 H_6 O}{1 \ mol \ C_2 H_4} = 3.626 \ mol \ C_2 H_6 O$$

$$55.19 \ g \ H_2 O * \frac{1 \ mol \ H_2 O}{18.02 \ g \ H_2 O} * \frac{1 \ mol \ C_2 H_6 O}{1 \ mol \ H_2 O} = 3.063 \ mol \ C_2 H_6 O$$

Thus, H₂O is the limiting reactant

$$3.063 \ mol \ C_2H_6O * \frac{46.07 \ g \ C_2H_6O}{1 \ mol \ C_2H_6O} * \frac{1 \ mL}{0.789 \ g} = 178.8 \ mL \ C_2H_6O$$

b. How much (mass) excess reactant remains?

55.19
$$g H_2 O * \frac{1 \, mol \, H_2 O}{18.02 \, g \, H_2 O} * \frac{1 \, mol \, C_2 H_4}{1 \, mol \, H_2 O} = 3.063 \, mol \, C_2 H_4 \, used \, up$$

101.7
$$g C_2 H_4 * \frac{1 \ mol \ C_2 H_4}{28.05 \ g \ C_2 H_4} = 3.626 \ mol \ C_2 H_4 \ initially$$

$$3.626\ mol - 3.063\ mol = 0.563\ mol\ C_2H_4\ remaining * \frac{28.05\ g}{1\ mol\ C_2H_4} = 15.8\ g\ C_2H_4\ remains$$

- 6. Complete and balance the following molecular equations, and then write the net ionic equation for each (note in past answer keys, we have written H⁺ in net ionic equations, but it is more correct to write H₃O⁺ instead. H⁺ doesn't actually exist itself in solution):
 - a. $HBr(aq) + NH_3(aq) \rightarrow$

$$HBr(aq) + NH_3(aq) \rightarrow Br^-(aq) + NH_4^+(aq)$$
 $H^+(aq) + Br^-(aq) + NH_3(aq) \rightarrow Br^-(aq) + NH_4^+(aq)$
 $H^+(aq) + NH_3(aq) \rightarrow NH_4^+(aq)$
 $H_3O^+(aq) + NH_2(aq) \rightarrow NH_4^+(aq) + H_2O(l)$

b. Aqueous hydrochloric acid and sodium acetate

Overall balanced:

$$HCl(aq) + NaC_2H_3O_2(aq) \rightarrow HC_2H_3O_2(aq) + NaCl(aq)$$

Total ionic:

$$H^{+}(aq) + Cl^{-}(aq) + Na^{+}(aq) + C_{2}H_{3}O_{2}^{-}(aq)$$

 $\rightarrow HC_{2}H_{3}O_{2}(aq) + Na^{+}(aq) + Cl^{-}(aq)$

Net ionic:

$$H^+(aq) + C_2H_3O_2^-(aq) \to HC_2H_3O_2(aq)$$

 $H_3O^+(aq) + C_2H_3O_2^-(aq) \to HC_2H_3O_2(aq) + H_2O(l)$

c. Aqueous perchloric acid and aqueous strontium hydroxide

Overall balanced:

$$2HClO_4(aq) + Sr(OH)_2(aq) \rightarrow Sr(ClO_4)_2(aq) + 2H_2O(l)$$

Total ionic:

$$2H^{+}(aq) + 2ClO_{4}^{-}(aq) + Sr^{2+}(aq) + 2OH^{-}(aq) \rightarrow Sr^{2+}(aq) + 2ClO_{4}^{2-}(aq) + 2H_{2}O(l)$$

Net ionic:

$$2H^+(aq) + 2OH^-(aq) \rightarrow 2H_2O(l)$$

$$2H_3O^+(aq) + 2OH^-(aq) \rightarrow 4H_2O(l)$$

The above equation can be simplified to:

$$H_3O^+(aq) + OH^-(aq) \rightarrow 2H_2O(l)$$

- 7. Starting with solid sucrose, C₁₂H₂₂O₁₁, describe how you would
 - a. Prepare 250 mL of a 0.250 M sucrose solution

$$0.250 M = \frac{0.250 \, mol \, C_{12} H_{22} O_{11}}{1 \, L} * 0.250 \, L = 0.0625 \, mol \, C_{12} H_{22} O_{11}$$

$$0.0625 \ mol \ C_{12}H_{22}O_{11} * \frac{342.3 \ g \ C_{12}H_{22}O_{11}}{1 \ mol \ C_{12}H_{22}O_{11}} = 21.39 \ g \ C_{12}H_{22}O_{11}$$

Measure out 21.39 g of sucrose, and add enough water to make 250 mL of solution.

b. Prepare 350.0 mL of 0.100 M $C_{12}H_{22}O_{11}$ starting with 3.00 L of 1.50 M $C_{12}H_{22}O_{11}$.

Determine how many moles of sucrose you need in the desired solution.

$$0.100 \ M \ C_{12}H_{22}O_{11} = \frac{0.100 \ mol \ C_{12}H_{22}O_{11}}{1 \ L} * 0.350 \ L = 0.035 \ mol \ C_{12}H_{22}O_{11}$$

You should check to make sure that you have at least this many moles of sucrose to begin with. Otherwise, you wouldn't have enough to prepare the desired solution.

$$1.50 \ M \ C_{12}H_{22}O_{11} = \frac{1.50 \ mol \ C_{12}H_{22}O_{11}}{1 \ L} * 3.00 \ L = 4.50 \ mol \ C_{12}H_{22}O_{11}$$

This confirms that we have enough moles of sucrose to make the desired solution. We need 0.035 mol of $C_{12}H_{22}O_{11}$.

$$0.035 \ mol \ C_{12}H_{22}O_{11} * \frac{1 \ L}{1.50 \ mol \ C_{12}H_{22}O_{11}} = 0.0233 \ L$$

So, we need 23.3 mL of the 1.50 M solution, and then the rest of the 350.0 mL volume should be water. The final solution volume is 350.0 mL.