#### AP CHEMISTRY LAB LIMITING REACTANT

#### Objectives:

- O To determine the limiting reactent in a selt mixture
- To determine the percent composition of each substance in a salt mixture.

#### Introduction.

Two factors limit the yield of products in a chemical reaction: (1) the amounts of starting materials (reactants) and (2) the percent yield of the reaction. Many experimental conditions, for example, temperature and pressure, can be adjusted to increase the yield, especially of a desired product, but because chemicals react according to fixed mole ratios (stoichiometrically), only a limited amount of product forms from given amounts of starting materials. The reactant determining the amount of product generated in a chemical reaction is called the **limiting reactant** in the chemical system.

To better understand the concept of the limiting reactant, Let's look at the reaction that is under investigation in this experiment. The reaction of sodium-phosphate dodecatrydrate, Ne.PO.12H<sub>2</sub>O and benium chloride dilrydrate, BeCl<sub>2</sub>2H<sub>2</sub>O. The molecular equation is:

2 Ne.PO:12H:O<sub>102</sub> + 3 BeCl:2H:O<sub>102</sub> -> Bec(PO:)<sub>201</sub> + 6 NeCl<sub>202</sub> + 30 H:O<sub>10</sub>

In this experiment an unknown self-mixture of Na,PO,12H,O and BaCl,2H,O is added to water producing insoluble Ba.(PO.). The mass of the imitial self-mixture and the mass of the final product, Ba.(PO.), are measured. A series of tests is also used to determine the limiting reactant. Finally, the percent composition of the original self-mixture is calculated.

#### Pre-laboratory Questions:

- The solubility of Be (PO): is 0.519 mg/L. How many milligrams and moles of Be (PO): dissolve in 200 ml of solution?
- Describe how the mass of Ba-(PO.): precipitate is determined in Part A.
- Describe how to test your unknown salt mixture for the presence of excess Na.PO.12H.O.
- 4. Why is fine porosity paper required for use in the filtering of the Ba<sub>2</sub>(PO<sub>3</sub>), precipitate?

Materials: Salt mixture, DI water, small besiders, scoop, electronic belance, ninse bottle, filter paper, framel, stir rod, watch glass, tongs, hot plate, 1000 mL besider, Erlenmeyer flask, dropper bottle 0.50 M Na.PO. 12H<sub>2</sub>O<sub>ing</sub>, dropper bottle of 0.50 M BaCl<sub>2</sub>2H<sub>2</sub>O<sub>ing</sub>

#### Precedure:

Two trials are recommended for this experiment. To hasten the analysis, weigh displicate unknown salt mixtures and simultaneously follow the procedure for each. Label the beakers accordingly for trial 1 and trial 2 to avoid the intermixing of samples and solutions.

#### A. Precipitation of Ba (PO.): from the salt mixture

- Weigh and Dissolve the Salt Mixture. Weigh (±0.001 g) on weighing paper about 1.0 gram of the unknown salt mixture.
  Record the mass for trial 1 in your laboratory notebook. Repeat for trial 2. Transfer the salt mixture to a labeled 400-ml beaker and add 200-ml of deionized water. Stir the mixture with a stirring rod for about 1 minute and then allow the precipitate to settle. Leave the stirring rod in the beaker.
- Digest the Precipitate. Cover the besker with a watchglass and maintain the solution hot (30-90°C) on a steam bath or with a low
  flame for 30 minutes. While the precipitate is being warmed, proceed to part A.3. Penodically check on the progress of heating.
  After 30 minutes, allow the precipitate to settle, it does not need to be cooled to room temperature.
- 3 Filter the Precipitate. Set up a gravity filtering apparatus. Weigh (+0.001 g) a piece of Whatman No.42 or Fisherbrand Q2 filter paper fine porosity filter paper. Fold it properly and seal it into the filter finnel with a small amount of decorated water. Discord the water in the receiving flask. Return to Part A.2.
- Separation of Supernatural in Part A.2. Once the supernatural has cleared in Part A.2, decant two 50-ml volumes into separate 100-ml beakers, labeled Beaker I and Beaker II. Save for part B.
- 5. Filter the Precipitate Salt. While the solution is still warm from Part A. 4, quantitatively transfer the precipitate to the filter funnel. Remove any precipitate on the wall of the beaker with a rubber policemen, and hot (\$0.90°C) water\*, wash the precipitate onto the filter. Rimse the Ba<sub>i</sub>(PO<sub>i</sub>), precipitate with two 5-ml portions of hot water (see Part A. 2).

  \*\*Ill hears a hot water buth up from for this purpose.
- 6. Dry and Weigh the Salt Precipitate. Remove the filter paper holding the precipitate from the filter funnel. Air-dry the precipitate on the filter paper until the next laboratory period or in a 110 °C constant-temperature drying oven overnight. Weigh (±0.001 g) the precipitate and filter paper. Record in notebook.

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