"<u>1.00 g of a Compound</u>" Lab Briefing

- > A *double replacement* reaction of two (*aq*) solutions will often produce a precipitate (*s*)
 - A: The *solid* precipitate can then be easily isolated through a *filtration process.*
 - B: Some ionic compounds contain *water molecules* (hydrates) that are trapped within their crystal structure and therefore must be *dried out*.
- LAB OBJECTIVE: To produce EXACTLY 1.00 grams of desired product (precipitate) using solubility rules of a double replacement reaction:
 - **<u>Pre-Lab</u>** Step 1: Write a *balanced chemical equation* for your given reaction.
 - <u>Pre-Lab</u> Step 2: Assign states of matter (aq or s) to <u>ALL</u> reactants and products in the chemical equation using *solubility rules*.
 - <u>Pre-Lab</u> Step 3: Calculate the <u>THEORETICAL MASS (g)</u> of <u>BOTH REACTANTS</u> needed to exactly produce the desired 1.00 grams of the *solid precipitate*.
 - A: 1.00 g of solid precipitate is the <u>GIVEN</u> → Use MASS → MASS stoichiometry to determine required mass of <u>BOTH REACTANTS</u>.
 Therefore, two (2) mass → mass stoichiometry calculation set-up required.
 - B: If ANY reactant is a <u>HYDRATE</u>, you <u>MUST</u> take the hydrate mass into account when determining its <u>total</u> MOLAR MASS.
 - C: Number of water molecules <u>MUST</u> also be written on the product side to correctly balance the equation.
 - Ex: MgSO₄ 7H₂O (aq) + Na₂CO₃ (aq) \rightarrow MgCO₃ (s) + Na₂SO₄ (aq)
 - $1 \text{ MgSO}_4 \bullet 7H_2O_{(aq)} + 1 \text{ Na}_2\text{CO}_{3 (aq)} \rightarrow 1 \text{ MgCO}_{3 (s)} + 1 \text{ Na}_2\text{SO}_{4 (aq)} + 7H_2O$
 - MgSO₄ 7H₂O → 120.37 g/mol MgSO₄ + 126.112 g/mol 7H₂O = 246.4822 g/mol
 - <u>Pre-Lab</u> Step 4: Watch the provided <u>VIDEO</u> for performing the proper filtration method/process.

- Actual Lab Step 1: Once the <u>THEORETICAL MASS</u> of both reactants are CALCULATED, dissolve both reactants in <u>SEPARATE</u> beakers using 25mL of <u>DISTILLED</u> water.
- Actual Lab Step 2: The two (2) DISSOLVED reactant solutions are then <u>MIXED</u> together in ONE of the two beakers (or into a third beaker) to produce the solid precipitate.
- Actual Lab Step 3: RECOVER precipitate through <u>FILTRATION</u> method using a piece of <u>PRE-MASSED filter paper</u>.
- Actual Lab Step 4: <u>CAREFULLY</u> remove <u>ALL</u> precipitate AND filter paper from funnel and place on <u>DRY</u> towel.
- Actual Lab Step 5: MASS dry precipitate AND filter paper together. <u>SUBTRACT</u> the total mass (precipitate + filter paper) from the original mass of filter paper to get just the mass of the precipitate.

AFTER Lab Step 1: Calculate % YIELD to determine how much of precipitate was <u>ACTUALLY</u> produced or recovered.

• How close is *your <u>ACTUAL</u>* mass to the 1.00 g <u>THEORETICAL</u> mass of precipitate?

% Yield = Actual Yield of Precipitate

----- x 100

Theoretical Yield of Precipitate

- AFTER Lab Step 2: Calculate % ERROR to determine how much of precipitate was NOT produced or recovered.
 - **% Error** = [Actual Theoretical]

----- x 100

Theoretical