CHAPTER 4
STOICHIOMETRY \& CHEMICAL EQUATIONS


## CHAPTER 6 INTRO TO CHEMICAL REACTIONS

## Writing and Balancing Chemical Equations [4.1]

¿WHY BALANCE CHEMICAL EQUATIONS?

$$
2 \mathrm{H}_{2}+\mathrm{O}_{2} \quad->\quad 2 \mathrm{H}_{2} \mathrm{O}
$$

(see "DrStephensonChemistry" youtube videos)

1. ID "lone" elements
(pure elements that are all "alone")
2. ID "twin" elements (element whose symbol appears once-and-only-once on each side of the equation. One of the twins lives on
the reactant-side, the other on the product-side)

3a. START with the Twin Element with the largest subscript.

3b. END by determining the coefficient of a Lone Element, if present.
4. Exploit opportunities to employ two tricks of the trade:
(i) cross-multiplication
(ii) fractionations

$$
\mathrm{CH}_{4}+\mathrm{O}_{2}->\quad \mathrm{H}_{2} \mathrm{O}
$$





CROSS MULTIPLY

$$
\frac{2}{1}\left(1 S^{K}+\frac{3}{2} C_{2} \rightarrow 1 \underline{S b}_{3}\right)
$$


'ONE-SHORT’ situation (2 or 3 known; 3 of 4 known; etc)


$$
\begin{aligned}
3+x & =2 y \\
x & =y \\
& \psi \\
3+x & =2 x \\
3 & =2 x-x \\
3 & =x
\end{aligned}
$$

## Classifying Chemical Reactions [4.2]

## Two General Classifications

(1) Ionic
(2) Oxidation-Reduction (REDOX)


IONIC reactions occur between ionic compounds
$\hookrightarrow$ recall, ionic compound is usually a Metal + Nonmetal
$\hookrightarrow$ or another way of putting it: a Type I or Type II compound
REDOX reactions are reactions between covalent compounds
$\hookrightarrow$ reactions between Type III compounds

Double Displacement Reactions

- Also known as a METATHESIS reaction
- Common vernacular is the PARTNER SWAP reaction
- Essentially, cations exchange their associated anions with each other
- Falls under the class of IONIC reactions
$\rightarrow$ lonic Compound = a Metal + a Nometal
$\hookrightarrow$ Ionic Compound = Type I or Type II compound


Predicting ionic reaction products

$$
A \hat{X}+B Y \rightarrow ? ?+? ? \quad A^{2 x} y_{2}^{-}+B X
$$

Given 2 Ionic Reactants: Procedure for Predicting Correct Molecular Equation
(1) Pair (elements)
(2) CHG neutral (compounds on Product-side)
(3) BCE (entire chemical equation)


Three Versions of Ionic Equations (each of which serves a different purpose)
(1) MOLECULAR (or Formula Unit) equation
(2) TOTAL IONIC equation
(3) NET IONIC equation


MOLECULAR >


NET IONIC >

$$
\begin{aligned}
& \mathrm{Pb}^{2+}+2 \mathrm{NO}_{3}^{-}+2 \mathrm{H}^{+}+\mathrm{SO}_{4}^{-} \rightarrow \mathrm{PbSO}_{4(\mathrm{~s})}+ \\
& \mathrm{Pb}^{2+}+\mathrm{SO}_{4}^{-} \rightarrow \mathrm{PbSO}_{4(\mathrm{~s})}
\end{aligned}
$$



## Equations for lonic Reactions

RECALL: ionic compounds are typically


- those composed of metals and non-metals (Type I \& II)
- Acids or Bases

Molecular:
$\mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}+\mathrm{H}_{2} \mathrm{SO}_{4} \rightarrow 2 \mathrm{HNO}_{3}+\mathrm{PbSO}_{4}$
Total lonic: $\mathrm{Pb}^{2+}+2 \mathrm{NO}_{3}{ }^{-}+2 \mathrm{H}^{+}+\mathrm{SO}_{4}{ }^{2-} \rightarrow 2 \mathrm{H}^{+}+2 \mathrm{NO}_{3}{ }^{-}+\mathrm{PbSO}_{4}(\mathrm{~s})$
Net lonic: $\mathrm{Pb}^{2+}+\mathrm{SO}_{4}{ }^{2-} \rightarrow \mathrm{PbSO}_{4}(\mathrm{~s})$

## SPECTATOR IONS

- cancel-out in Net Ionic
- are neither physically nor chemically changed by the rxn


(EX) Determination of Net Ionic Reaction
¿What is the net ionic equation for the following unbalanced equation?

$$
\mathrm{HCl}+\mathrm{Ca}(\mathrm{OH})_{2} \rightarrow \mathrm{CaCl}_{2}+\mathrm{H}_{2} \mathrm{O}
$$

$$
\begin{array}{rlr}
2 \mathrm{HCl}+{\mathrm{Ca}(\mathrm{OH})_{2}}_{\substack{\times \\
\downarrow}} \rightarrow \mathrm{CaCl}_{2}+2 \mathrm{H}_{2} \mathrm{O} & \text { Molecular } \\
2 \mathrm{H}^{+}+2 \mathrm{Cl}^{-}+\mathrm{Ca}_{2}^{2+}+2 \mathrm{HO}^{-} & \rightarrow \mathrm{Ca}^{2+}+2 \mathrm{Cl}^{-}+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) & \text { Total Ionic } \\
2 \mathrm{H}^{+}+2 \mathrm{HO}^{-} & \longrightarrow 2 \mathrm{H}_{2} \mathrm{O}(\mathrm{ll}) & \text { Net lon :c }
\end{array}
$$

## Precipitation Reactions



## SOLUBLE

- Group 1 cations ( $\mathrm{Ll}^{+}-\mathrm{CS}^{+}$)
- ammonium $\mathrm{NH}_{4}^{+}$
- halides $\mathrm{Cl}^{-}, \mathrm{Br}^{-}, \mathrm{I}^{-} \longrightarrow$ EXCEPT halides of $\mathrm{Ag}^{+}, \mathrm{Hg}_{2}{ }^{2+}, \mathrm{Pb}^{2+}$
- acetate $\mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{-}$
- bicarbonate $\mathrm{HCO}_{3}^{-}$
- nitrate $\mathrm{NO}_{3}{ }^{-}$
- chlorate $\mathrm{ClO}_{3}^{-}$
- sulfate $\mathrm{SO}_{4}{ }^{2-} \longrightarrow$ EXCEPT sulfates of $\mathrm{Ag}^{+}, \mathrm{Hg}_{2}{ }^{2+}, \mathrm{Pb}^{2+} \mathrm{Ba}^{2+}, \mathrm{Ca}^{2+}, \mathrm{Sr}^{2}$

IN-SOLUBLE $\rightarrow$ EXCEPT in presence of Group 1 cations and Ammonia - carbonate $\mathrm{CO}_{3}{ }^{2-}$

- chromate $\mathrm{CrO}_{4}{ }^{2-}$
- phosphate $\mathrm{PO}_{4}{ }^{3-}$
- sulfide $\mathrm{S}^{2-}$
- hydroxide $\mathrm{OH}^{-} \longrightarrow$ EXCEPT hydroxides of Group 1 cations and $\mathrm{Ba}^{2+}$


## Precipitation Reactions and Solubility Rules

- PRECIPITATION RXN - substance reacts to form solid product
- Examples:
$\hookrightarrow$ KIDNEY STONES (several varieties)

$$
\mathrm{Ca}^{2+}+\mathrm{C}_{2} \mathrm{O}_{4}{ }^{2-} \underset{\text { culcalm oxilate }}{\mathrm{CaC}_{2} \mathrm{O}_{4}} \text { K } \mathrm{K}_{\mathrm{sp}}=\left[\mathrm{Ca}^{2+}\right]\left[\mathrm{C}_{2} \mathrm{O}_{4}^{2-}\right]
$$

$\rightarrow$ CORAL REEF: CaCO3 + sea salt + algae for color

- SOLUBILITY / INSOLUBILITY
$\hookrightarrow$ are condition dependent (solube under circumstances, not others)... but...
$\longrightarrow$ often, cast solubility as simply "Yes/No" or "is soluble / is insoluble"

Solubilities of Common Ionic Compounds in Water

| Soluble compounds contain <br> - group 1 metal cations ( $\mathrm{Li}^{+}, \mathrm{Na}^{+}, \mathrm{K}^{+}, \mathrm{Rb}^{+}$, and $\mathrm{Cs}^{+}$) and ammonium ion $\left(\mathrm{NH}_{4}{ }^{+}\right)$ <br> - the halide ions $\left(\mathrm{Cl}^{-}, \mathrm{Br}^{-}\right.$, and $\left.\mathrm{I}^{-}\right)$ <br> - the acetate $\left(\mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}{ }^{-}\right)$, bicarbonate $\left(\mathrm{HCO}_{3}{ }^{-}\right)$, nitrate $\left(\mathrm{NO}_{3}{ }^{-}\right)$, and chlorate $\left(\mathrm{ClO}_{3}{ }^{-}\right)$ions <br> - the sulfate $\left(\mathrm{SO}_{4}{ }^{-}\right)$ion | Exceptions to these solubility rules include <br> - halides of $\mathrm{Ag}^{+}, \mathrm{Hg}_{2}{ }^{2+}$, and $\mathrm{Pb}^{2+}$ <br> - sulfates of $\mathrm{Ag}^{+}, \mathrm{Ba}^{2+}, \mathrm{Ca}^{2+}, \mathrm{Hg}_{2}{ }^{2+}$, $\mathrm{Pb}^{2+}$, and $\mathrm{Sr}^{2+}$ |
| :---: | :---: |
| Insoluble compounds contain <br> - carbonate $\left(\mathrm{CO}_{3}{ }^{2-}\right)$, chromate $\left(\mathrm{CrO}_{4}{ }^{2-}\right)$, phosphate $\left(\mathrm{PO}_{4}{ }^{3-}\right)$, and sulfide $\left(\mathrm{S}^{2-}\right)$ ions <br> - hydroxide ion $\left(\mathrm{OH}^{-}\right)$ | Exceptions to these insolubility rules include <br> - compounds of these anions with group <br> 1 metal cations and ammonium ion <br> - hydroxides of group 1 metal cations and $\mathrm{Ba}^{2+}$ |

Table 4.1

| 10r |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| sodium ion | $\mathrm{Na}^{+}$ | $\checkmark$ |  |  |
| potassium ion | $\mathrm{K}^{+}$ | $\checkmark$ |  |  |
| ammonium | $\mathrm{NH}_{4}^{+}$ | $\checkmark$ |  |  |
| nitrate | $\mathrm{NO}_{3}{ }^{-}$ | $\checkmark$ |  |  |
| acetate | $\mathrm{AcO}^{-}$ | $\checkmark$ |  |  |
| halogen ion | $\mathrm{Cl}^{\prime}, \mathrm{Br}^{-} \mathrm{l}^{-}$ | $\checkmark$ |  | salts with... $\mathrm{Ag}^{+}, \mathrm{Hg}_{2}{ }^{2+}, \mathrm{Pb}^{2+}$ |
| sulfate | $\mathrm{SO}_{4}{ }^{2-}$ | $\checkmark$ |  | salts with... $\mathrm{Ba}^{2+}, \mathrm{Ca}^{2+}, \mathrm{Pb}^{2+}, \mathrm{Hg}^{2+}$ |
| sulfide | $\mathrm{S}^{2-}$ |  | $\checkmark$ | salts with... $\mathrm{Na}^{+}, \mathrm{K}^{+}, \mathrm{NH}_{4}^{+}, \mathrm{Mg}_{2}{ }^{+}, \mathrm{Ca}_{2}{ }^{+}$ |
| carbonate | $\mathrm{CO}_{3}{ }^{2-}$ |  | $\checkmark$ | salts with... $\mathrm{Na}^{+}, \mathrm{K}^{+}, \mathrm{NH}_{4}^{+}$ |
| phosphate | $\mathrm{PO}_{4}{ }^{3-}$ |  | $\checkmark$ | salts with... $\mathrm{Na}^{+}, \mathrm{K}^{+}, \mathrm{NH}_{4}^{+}$ |
| arsenate | $\mathrm{AsO}_{4}{ }^{3-}$ |  | $\checkmark$ | salts with... $\mathrm{Na}^{+}, \mathrm{K}^{+}, \mathrm{NH}_{4}^{+}$ |
| oxide | $\mathrm{O}^{2-}$ |  | $\checkmark$ | salts with... Group 1 \& 2 metals |
| hydroxide | HO |  | $\checkmark$ | strong bases |

(EX) Predict PPT Reactions
¿(a) Which solution could be used to precipitate the barium ion, BaL+, in a water sample: sodium chloride, sodium hydroxide, or sodium sulfate? (b) What is the formula for the expected precipitate?
(i) $\mathrm{Ba}^{2+}+\mathrm{NaCl}_{\mathrm{a}} \rightarrow \mathrm{BaCl}_{2} \longleftarrow$ solute
(ii) $\mathrm{Ba}^{2+}+\mathrm{NaOH} \longrightarrow \mathrm{Ba}_{\mathrm{a}}(\mathrm{OH})_{2} \longleftarrow$ soluble
(iii) $\mathrm{Ba}^{2+}+\mathrm{NaSO}_{4} \rightarrow \mathrm{BaSO}_{4} \leftharpoonup$ precip.taAes

- ACID-BASE REACTION
$\rightarrow$ hydrogen ion (aka proton), $\mathrm{H}^{+}$, is transferred, or
$\hookrightarrow$ hydronium ion, $\mathrm{H}_{3} \mathrm{O}^{+}$, is transferred
- ARRHENIUS ACID - donates $\mathrm{H}^{+}$in water
- ARRHENIUS BASE - donates hydroxide ion, $\mathrm{OH}^{-}$(aka $\mathrm{HO}^{-}$), in water

Strong vs. Weak Acids
$\rightarrow$ STRONG ACID - dissociates 100\%
$\longrightarrow$ WEAK ACID - dissociates <100\%

$$
\begin{aligned}
H A & \rightarrow A^{-}+H^{+} \\
\mathrm{BOH} & \rightarrow \mathrm{~B}^{+}+\mathrm{HO}^{-}
\end{aligned}
$$

$$
\begin{aligned}
& \begin{array}{ccc}
\mathrm{HCl} & \mathrm{H}^{+}+\mathrm{Cl}^{-} \\
100 & \varnothing & \varnothing \\
\varnothing & 100 \quad 100
\end{array}
\end{aligned}
$$


eg: $\mathrm{HCl}+\mathrm{NaOH} \rightarrow \mathrm{NaCl}+\mathrm{H}_{2} \mathrm{O}$

Acid-Base Reaction is a type of Double Displacement

$$
\begin{array}{lrl}
\text { Molecular: } & \mathrm{Ba}(\mathrm{OH})_{2}+2 \mathrm{HNO}_{3} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}_{(l)^{+}} \mathrm{Ba}\left(\mathrm{NO}_{3}\right)_{2} \\
\text { Total lonic: } & \mathrm{Ba} a^{2+}+2 \mathrm{HO}^{-}+2 \mathrm{H}^{+}+2 \mathrm{NO}_{3}^{-} \rightarrow & 2 \mathrm{H}_{2} \mathrm{O}()^{+} \mathrm{Ba}^{2+}+2 \mathrm{NO}_{3}^{-} \\
\text {Net lonic: } & 2 \mathrm{HO}^{-}+2 \mathrm{H}^{+} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}
\end{array}
$$

Oxidation-Reduction Reactions (buckle up... here's where it starts to get hairy)


Definition: REDOX

- OXIDATION = 个 O.N.
- REDUCTION $=\downarrow$ ON.

OIL $\{$ election flow $(-)$ is opposite the RIG direction of OP( + )
"AGENTS" (facilitates an action): What it is, not what it does

- Oxidizing agent (OXIDANT) = species that causes something else to become oxidized ... it is, itself, reduced.
- Reducing agent (REDUCTANT) = species that causes something else to become reduced... it is, itself, oxidized.


## Oxidation Numbers

oxidation number (oxidation state) - charge atom would bear if it were part of an ionic compound.

## Rules for Assigning ON's

1. The oxidation number of an atom in an elemental substance is zero.
2. The oxidation number of a monatomic ion is equal to the ion's charge.
3. Oxidation numbers for common nonmetals are usually assigned as follows:

- Hydrogen: +1 when combined with nonmetals, -1 when combined with metals
- Oxygen: -2 in most compounds, sometimes -1 (so-called peroxides, O 2 2- ),
very rarely - 1 (so-2 called superoxides, $\mathrm{O} 2-$ ), positive values when combined with $F$ (values vary)
- Halogens: -1 for $F$ always, -1 for other halogens except when combined with oxygen or other halogens (positive oxidation numbers in these cases, varying values) 4. The sum of oxidation numbers for all atoms in a molecule or polyatomic ion equals the charge on the molecule or ion.

5. GROUP $1=+1$
6. GROUP $2=+2$
7. GROUP 3 = usually +3
8. Transition metals - Type I / Type II rules as guidance
9. Polyatomic lons considered as SINGLE GROUP.
(EX) Assign ON's. [4.5b]
¿Assign oxidation states to the elements whose atoms are underlined in each of the following compounds or ions:
(a)

(C)
(b)

(d)


## COMBUSTION REACTION (Redox subclass)

- Reaction of fuel
- ex: hydrocarbon (HC) + oxidant (esp. $\mathrm{O}_{2}$ )

4 Types of "REDOX questions. (easier to harder)
(1) Is this reaction a redox?
(2) What is oxidized? (or, What is reduced?)
(3) What is oxidizing agent? (or, What is reducing agent?)
(4) How many electrons are transferred
<see example problem next page>
(EX) I.D. Redox Combustion Run Components
¿Consider the formation of a rust from a pure iron, according to the following equation. What is...
(a) the material that is reduced?; (b) the material that is oxidized?; (c) how many electrons were transferred?; (d) the oxidizing agent?; (e) the reducing agent?

$$
4 \mathrm{Fe}+3 \mathrm{O}_{2} \rightarrow 2 \mathrm{Fe}_{2} \mathrm{O}_{3}
$$


(EX) I.D. Redox Combustion Run Components
¿Consider the formation of a rust from a pure iron, according to the following equation. What is... (a) the material that is reduced?; (b) the material that is oxidized?; (c) how many electrons were transferred?; (d) the oxidizing agent?; (e) the reducing agent?


TIP: any reaction which forms a compound from an element (or vice versa) is necessarily a REDOX reaction

Single-Displacement (Replacement) Reaction: another REDOX subclass:
$\longrightarrow$ only 1 moitey displaced
$\hookrightarrow$ especially common reactions with METALS


Balancing Redox Reactions via the Half-Reaction Method

$$
\text { (page } 189 / 196 \text { ) }
$$

<'BALANCING REDOX EQUATIONS' NOT COVERED ON THIS EXAM>
Notes-Part Ka"

Reaction Stoichiometry [4.3]

Review from Last Chapter


(EX) 'gram-to-gram' Stoichiometry Problem
¿ For the synthesis of water from hydrogen and oxygen, how many grams of hydrogen are required to react with 128 g of oxygen,?


$$
2 \mathrm{H}_{2}+\mathrm{O}_{2}->
$$



Hin $\mathrm{gH}_{2}=\frac{128 \mathrm{gO}}{2} \cdot \frac{1 \mathrm{molO}_{2}}{32 \mathrm{gO}_{2}} \cdot \frac{2 \mathrm{molH}_{2}}{1 \mathrm{molO}_{2}} \cdot \frac{2 \mathrm{gH}_{2}}{1 \mathrm{molH}_{2}}=16 \mathrm{gH}_{2}$
( $\mathrm{gH}_{2} \mathrm{O}=\frac{128 \mathrm{gO}_{2}}{} \cdot \frac{1 \mathrm{molO}_{2}}{32 \mathrm{gO}} \cdot \frac{2 \mathrm{molH}_{2} \mathrm{O}}{1 \mathrm{molO}_{2}} \cdot \frac{18 \mathrm{gH}_{2} \mathrm{O}}{1 \mathrm{molH}_{2} \mathrm{O}}=144 \mathrm{gH}_{2} \mathrm{O}$

(EX) $g(a) \rightarrow g(b) \quad$ [4.11b]
¿What mas of CO is required to react with 25.13 g Fe 2 O 3 according to the equation:


DA
25.13 g
$\square \mathrm{g}$

$$
\mathrm{Fe}_{2} \mathrm{O}_{3}+3 \mathrm{CO} \rightarrow 2 \mathrm{Fe}+3 \mathrm{CO}_{2}
$$


(EX) ga) $->\mathrm{g}(\mathrm{b})$
[4.11b]
¿What mass of SO2 contains the same mass of oxygen as is contained in 33.7 g of As 2O5?

$? \frac{\mathrm{~g} \cdot \mathrm{SO}_{2}}{}=\frac{33.7 \mathrm{~g} \cdot A s_{2} \mathrm{O}_{5}}{1} \cdot \frac{1 \mathrm{~mol} \cdot \mathrm{As}_{2} \mathrm{O}_{5}}{229.8 \mathrm{~g} \cdot \mathrm{As}_{2} \mathrm{O}_{5}} \cdot \frac{5 \mathrm{~mol} \cdot \mathrm{O}}{1 \mathrm{~mol} \cdot \mathrm{As}_{2} \mathrm{O}_{5}} \cdot \frac{1 \mathrm{~mol} \cdot \mathrm{SO}_{2}}{2 \mathrm{~mol} \cdot \mathrm{O}} \cdot \frac{64.1 \mathrm{~g} \cdot \mathrm{SO}_{2}}{1 \mathrm{~mol} \cdot \mathrm{SO}_{2}}=23.5 \mathrm{~g} \cdot \mathrm{~S}^{\prime}$

Limiting Reactant

$$
\begin{gathered}
2 \text { bread } \\
14
\end{gathered}+\underset{4}{1 \text { meat } \rightarrow} \begin{gathered}
1 \text { sandwich } \\
? ? ? \\
7
\end{gathered}
$$

CLAIM: with 14 slices of bread and 4 slices of meat, I can make 7 sandwiches

ARGUMENT:

$$
? ? \text { sand }=\frac{14 \text { bread }}{1 \text { sandwich }} \frac{2 \text { bread }}{2 \text { sandwiches }}
$$

COUNTER
ARGUMENT: $\quad ? ?$ sand $=\frac{4 \text { meat }}{1 \text { sandwich }} \frac{1 \text { meat }}{1 \text { sandwiches }}$
ANSWER: cannot make more product than the material in least supply (Limiting Reagent) will allow.

(EX) ID Limiting Reagent

$$
\begin{array}{lc}
38.0 \mathrm{~g} & 225 \mathrm{~g} \\
2 \mathrm{H}_{2}+\mathrm{O}_{2} \rightarrow> & \square \mathrm{g} \\
2 \mathrm{H}_{2} \mathrm{O}
\end{array}
$$



$$
2 \mathrm{H}_{2}+\mathrm{O}_{2} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}
$$

$$
? ? \mathrm{gW}=\frac{38.0 \mathrm{gH}}{2} \cdot \frac{1 \mathrm{molH}_{2}}{2.02 \mathrm{gH}_{2}} \cdot \frac{2 \mathrm{molH}_{2} \mathrm{O}}{2 \mathrm{molH}_{2}} \cdot \frac{18.0 \mathrm{gH}_{2} \mathrm{O}}{1 \mathrm{molH}_{2} \mathrm{O}}=339 \mathrm{gH}_{2} \mathrm{O}
$$

$$
? ? \mathrm{gW}=\frac{225 \mathrm{gO}_{2}}{} \cdot \frac{1 \mathrm{molO}_{2}}{32.00 \mathrm{gO}_{2}} \cdot \frac{2 \mathrm{molH}_{2} \mathrm{O}}{1 \mathrm{molO}_{2}} \cdot \frac{18.0 \mathrm{gH}_{2} \mathrm{O}}{1 \mathrm{molH}_{2} \mathrm{O}}=253 \mathrm{gH}_{2} \mathrm{O} \quad \text { LIIITING }
$$

(EX) Calculate Excess of Excess Reagent
¿EXCESS: back-catculate to find the am. of Excess Reagent needed to make the lesser amt of product, then subtract/from the amt. of Excess Reagent available at the start?

Buck -C

$? ? \mathrm{gH}_{2}=\frac{253 \mathrm{gH}_{2} \mathrm{O}}{1 \mathrm{molH}_{2} \mathrm{O}} 18.0 \mathrm{gH}_{2} \mathrm{O} \cdot \frac{2 \mathrm{molH}_{2}}{2 \mathrm{molH}_{2} \mathrm{O}} \cdot \frac{2.02 \mathrm{gH}_{2}}{1 \mathrm{molH}_{2}}=28.4 \mathrm{gH}_{2}$ dit
?? $\mathrm{g} \mathrm{H}_{2}($ excess $)=38.0-28.4=9.6 \mathrm{gH}$

$$
\text { \%Yield }=\frac{\text { actual } \text { ( } \text { measured })}{\text { theoretical }(\text { calculated })} \stackrel{100}{\leftarrow} \text { So.dintic calc }
$$

(EX) Calculate \% Yield from grams of reactant ¿What is the \% yield of a rxn that produces 12.5 g of the gas Freon from 32.9 g of CCI4 in excess HF?
<see answer next page>

## LECTURE STOP

(EX) Calculate \% Yield from grams of reactant ¿What is the \% yield of a rxn that produces 12.5 g of the gas Freon from 32.9 g of CCI4 in excess HF?

$$
25.9_{2} \text { (ther) }
$$

$$
32.9 \mathrm{~g} \quad \text { excess } \quad 12.5 \mathrm{~g} \text { (measured) }
$$

$$
\mathrm{CCl}_{4}+2 \mathrm{HF} \rightarrow \mathrm{CF}_{2} \mathrm{Cl}_{2}+2 \mathrm{HCL}
$$

(1) $\%$ Yield $=\frac{12.5 \mathrm{~g}}{\text { ??? theoretical }} \cdot 100$
(2) ${ }^{2} ? g_{\mathrm{CF}_{2} \mathrm{Cl}_{2}}=\frac{32.9 \mathrm{gCCl}}{4} \cdot \frac{1 \mathrm{molCCl}_{4}}{153.82 g \mathrm{gCl}_{4}} \cdot \frac{1 \mathrm{molCF}_{2} \mathrm{Cl}_{2}}{1 \mathrm{molCl}_{4}} \cdot \frac{120.91 \mathrm{gCF} \mathrm{Cl}_{2}}{1 \mathrm{molCF}_{2} \mathrm{Cl}_{2}}=25.9 \mathrm{gCF}_{2} \mathrm{Cl}_{2}$
(3) $\%$ Yield $=\frac{12.5 g}{25.9 g} \cdot 100=48.3 \%$

(EX) Calculate \% Yield from given amount of product
¿If you ran this reaction with 128 g of oxygen, but you only recovered 105 g of water, how well did you do?


$$
\left.\begin{array}{c}
\begin{array}{c}
16.0 \mathrm{~g} \quad 128 \mathrm{~g} \\
2 \mathrm{H}_{2}+\mathrm{O}_{2} \rightarrow
\end{array} \begin{array}{c}
105 \mathrm{~g} \\
144 \mathrm{~g}
\end{array} \\
2 \mathrm{H}_{2} \mathrm{O}
\end{array}\right] \begin{gathered}
\text { part } \\
\text { \% Yield }=\frac{105}{\text { whole }} \cdot 100=0.6935 \cdot 100=69.3 \% \\
144
\end{gathered} 100=0 .
$$

QUANTITATIVE ANALYSIS - det' $n$ of amt of material present.
(1) Titration,
(2) Gravimetric, \&
(3) Combustion

## Titration

DESCRIPTION: a known amount of Titrant (known conc) is added to a known amount of Analyte (unknown conc) until th Equivalence Point (point at which the amount of Titrant and Analyse are equal, per Indicator at the End Point).



Figure 1: Titration Setup

$$
M_{A} V_{A}=M_{B} V_{B}
$$

(EX) Calculate Molarity and g(Titrant used)
¿ 50.0 mL of HCl is titrated to the equivalence point with 23.4 mL of 0.216 M NaOH . What is the molarity of the HCl solution?

$$
\mathrm{HCl}+\mathrm{NaOH} \rightarrow \mathrm{NaCl}+\mathrm{H}_{2} \mathrm{O}
$$



$$
\begin{aligned}
\mathrm{M} \cdot \mathrm{~V} & =\mathrm{M}^{\prime} \cdot \mathrm{V}^{\prime} \\
50.0 \cdot \mathrm{X} & =23.4 \cdot 0.216 \\
\mathrm{X} & =0.101 \mathrm{M}
\end{aligned}
$$


(b) ¿How many grams of HCl were in the titrated sample?
$\begin{array}{ll}50.0 \mathrm{~mL} & 23.4 \mathrm{~mL} \\ & 0.216 \mathrm{~mol} / \mathrm{L}\end{array}$
$\mathrm{HCl}+\mathrm{NaOH} \rightarrow \mathrm{NaCl}+\mathrm{H}_{2} \mathrm{O}$

$$
? ? g_{a}=\frac{23.4 m L_{b}}{0.216 \mathrm{~mol}_{b}} \cdot \frac{1 \mathrm{~mol}_{a}}{1000 m L_{b}} \cdot \frac{36.46 g_{a}}{1 m o l_{b}}=0.184 g_{a}
$$

## Gravimetric Analysis

$\hookrightarrow$ based on CHANGE IN MASS (which affects the pull of Gravity)
$\hookrightarrow$ ex: dehydration
WATER LOSS = mass(before) - mass(after)

(a) $? \frac{g_{C I}}{1}=\frac{1.0881 g_{A g C l}}{1} \cdot \frac{1 \mathrm{~mol}_{A g C l}}{143.32 g_{A g C l}} \cdot \frac{1 \mathrm{~mol}_{c r}}{1 \mathrm{~mol}_{A g C l}} \cdot \frac{35.45 g_{c r}}{1 \mathrm{~mol}_{c r}}=0.2691 g_{c r}$
(b) $\% \mathrm{Cl}=\frac{0.2691 \mathrm{~g}}{1.1324} \cdot 100=23.76 \%$

Combustion Analysis

- Oxidizes hydrocarbon (reacts with O 2 ) to blast material into pieces, then in captures those H 2 O and CO2 pieces.
$\rightarrow \mathrm{H} 2 \mathrm{O}$ gives measure of H
$\hookrightarrow \mathrm{CO2}$ given measure of C

$$
2 \mathrm{C}_{2} \mathrm{H}_{6}+7 \mathrm{O}_{2} \rightarrow 4 \mathrm{CO}_{2}+6 \mathrm{H}_{2} \mathrm{O}
$$


(EX) Calculate Emp. Formula from Combustion Data [4.16b] ¿A 0.00215-g sample of polystyrene, a polymer composed of carbon and hydrogen, produced 0.00726 g of CO 2 and 0.00148 g of H 2 O in a combustion analysis. What is the empirical formula for polystyrene?

$$
\mathrm{C}_{x} \mathrm{H}_{y} \leftarrow \text { TARGET }
$$




(ii) $\square \mathrm{mol} H=$| $0.00148 \mathrm{~g} \omega$ | $1 \mathrm{~mol} \omega$ | $2 \mathrm{~mol} H$ |
| :--- | :--- | :--- |
|  | $18.02 \mathrm{~g} \omega$ | $1 \mathrm{~mol} \omega$ |$=1.64 \varepsilon-4 \mathrm{mdl} \mathrm{H}$

(iii) $C_{x} H_{y} \Rightarrow C_{\frac{1.65 \varepsilon-4}{1.64 \varepsilon-4}} H_{\frac{1.64 \varepsilon-4}{1.64 \varepsilon-4}} \Rightarrow C_{1} H_{1} \Rightarrow C H$

