

## Q) Weight



1) Density =  $\frac{\text{Mass}}{\text{Volume}} \approx \frac{\text{g}}{\text{mL/cm}^3}$

2) \* Mass of water lost upon heating  
= Mass of alum - Mass of anhydrous

$$\rightarrow \text{Moles} = \frac{\text{Mass}}{\text{Molar Mass}}$$

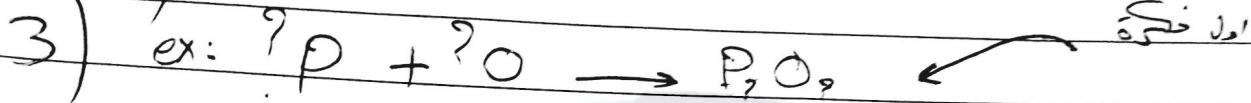
\* percentage of water of crystallizing by Mass  
= Mass of lost upon heating  $\times 100\%$

$$\rightarrow \text{value of "X"} = \frac{\text{Moles of water}}{\text{Moles of anhydrous salt}}$$

$$\rightarrow \frac{(\text{Mass of alum} - \text{Mass of water})}{\text{Molar mass of anhydrous}}$$

$$\frac{(\text{Mass of alum} - \text{Mass of anhydrous})}{\text{Molar mass of water}}$$

A)



Mass of white oxide (P<sub>2</sub>O<sub>5</sub>) - Mass of (P)  
= Mass of (O<sub>2</sub>)

→ Mass = moles of O<sub>2</sub> and P  
M/M

empirical formula → P<sub>n</sub>O<sub>n</sub>

division by smallest number of moles

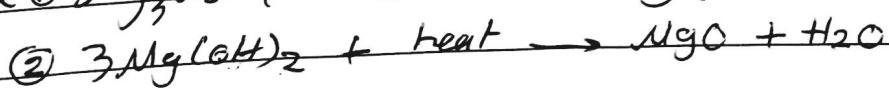
→ mass percent of "X" in X<sub>0</sub> (oxide)  
$$-\frac{\text{mass } X}{\text{mass } X_0} \times 100\% \Rightarrow \text{experimentally } \underline{\underline{x_1}}$$

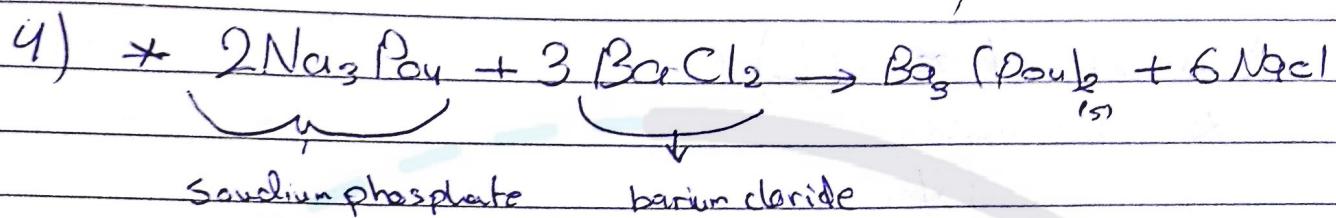
~~percent error~~

Mass percent of Mg in MgO (calculated)

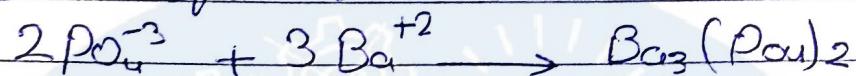
= Molar Mass of Mg × 100% x<sub>2</sub>  
Molar Mass of MgO

~~Mass~~  
→ percentage error =  $\left[ \frac{|x_2 - x_1|}{x_2} \right] \times 100\%$





\* net Ionic equation :-



→ \* if we have moles → moles  
 "moles" limiting reactant  $\rightarrow$  المدخل المحدود

→ moles of product  $\xrightarrow{\text{परिवर्तनी}}$   
 $(\text{मात्रा}) \Rightarrow \text{जलालिम}$

$$* \text{ Mass of excess} = \underline{\underline{M_{\alpha}}}$$

$$\text{Mass of salt mix} - (\text{Mass of limiting} + \text{Mass of Ba--})$$

$$* \text{ Mass percentage of limiting} = \frac{\text{Mass of limiting}}{\text{Mass of salt mix}} \times 100\%$$

\* test for excess of  $P_{O_2}$

we add 2 drop of BaCl<sub>2</sub> solution to beaker if precipitate forms then PO<sub>4</sub><sup>3-</sup> excess ba limiting

\* test for excess back  
S31(wi)

$$5) * n = M \times V(L)$$

\*  $n = \text{mass}$

$$\frac{\text{mass}}{M.M}$$

→ Number of moles of base = Number of moles of acid

\* if the Indicator phenolphthalein, which is colorless in acid and pink in base

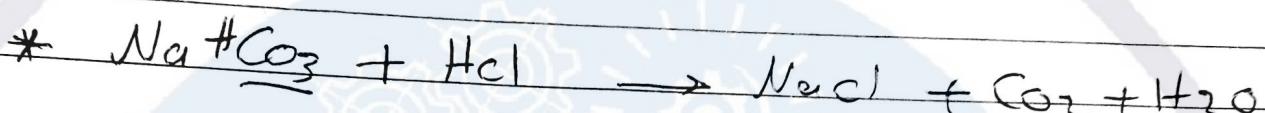
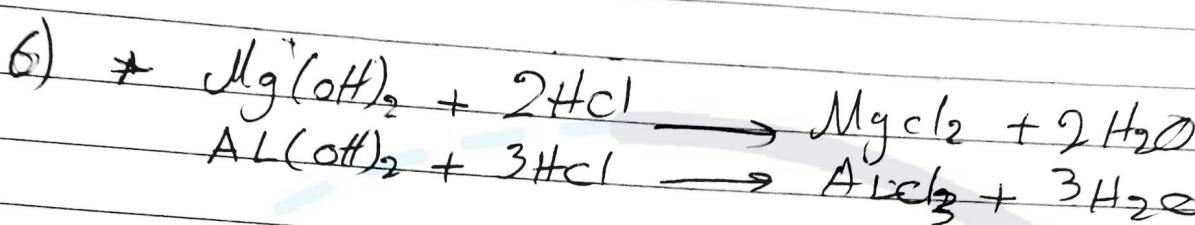
\* we have volume and we have molarity → moles of base

Molarity ← moles of acid  
mass → 1:1

\* Mass percent acetic acid in Vinegar

$$= \frac{\cancel{\text{mass}} \times \text{Molarity}}{10 \times \text{density}} \%$$

Δ 15° ✓  
1.06 g/cm³  
500 ml



\* Mass of antacid sample

~ 9

HCl

NaOH

$$n_1 = V \times N$$

$$\frac{n}{2} = V \times N$$

used to dissolve buret reading  
antacid.

same HCl

~~Excess HCl~~  $\Delta V$

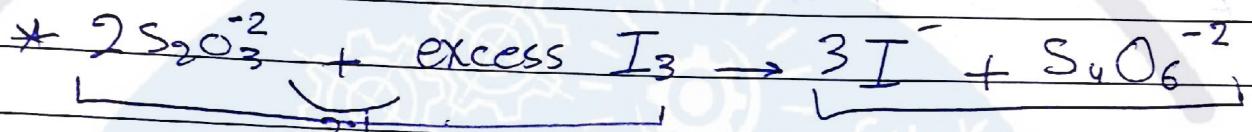
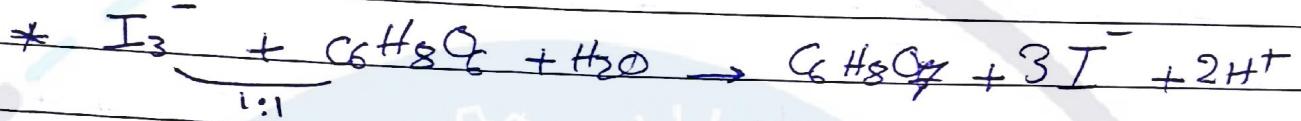
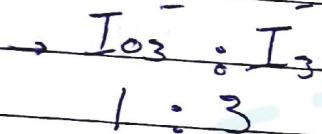
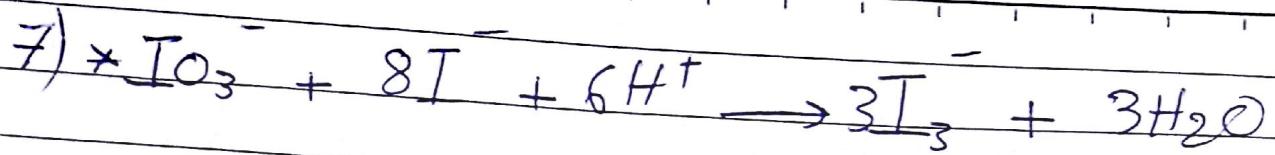
$n_{\text{excess of HCl}} = n_2$

$$n_{\text{nut}} = n_1 - n_2$$

needed to neutralize  
antacid tablet

\* Neutralizing capacity  
of antacid

$$= \frac{n_{\text{nut}}}{\text{Mass of antacid}}$$



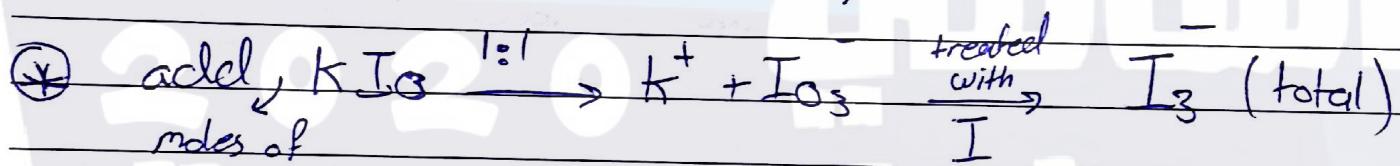
deep blue

colorless

mass for sample

1:3

1(x3)



$\oplus \text{ } *n \text{ of } \text{Na}_2\text{S}_2\text{O}_3 = N \text{ } *V(L)$

, 14)

→ buret (F-I)

\* n of excess  $I_3^-$  ( $\div 2$ )

~~Mass of  $S_2O_3$~~

121

(+) ~~n~~ of  $I_3^-$  (Total - excess)

reduced by  
 $\text{C}_6\text{H}_{5}\text{O}_5$

Mass  
of

x 111

\* n of  $C_6H_8O_6$  <

15

Mass percent

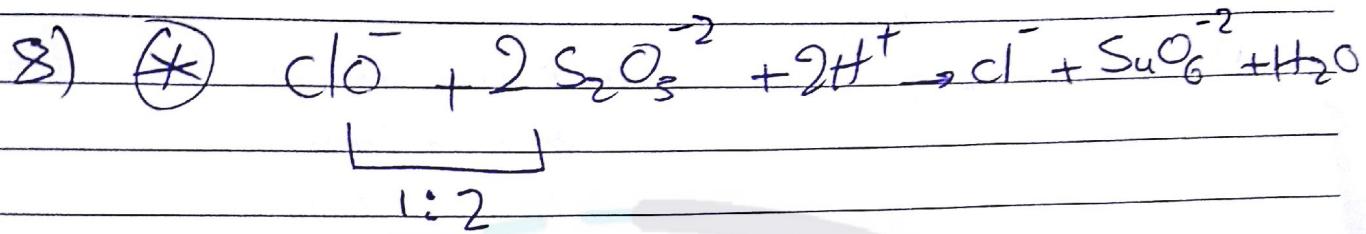
$\rightarrow \frac{\text{mass}}{\text{sample}} \times 100\%$

sample -?

red brown → pale yellow → deep blue → light greenish-yellow

$\text{Fe}_2\text{O}_3$        $\text{S}_2\text{O}_3^{2-}$ ?      starch       $\text{S}_2\text{O}_3^{2-}$       ?

Five Apple  
-G-



~~\*~~ Volume of bleach solution

~~\*~~ Volume of  $\text{Na}_2\text{S}_2\text{O}_3$  solution (buret reading (I - I))

$$\cancel{\ast} \text{ n of } \text{ClO}^- \text{ reduced} \xleftarrow[\text{(1:2)}]{\div 2} \cancel{\ast} \text{ n of } \text{Na}_2\text{S}_2\text{O}_3$$

$\frac{\text{volume}}{\text{of}} \text{ bleach}$

Molarity of diluted bleach

~~\*~~ diluted factor

Molarity of original bleach

$$\cancel{\ast} \text{ Mass percent} = \text{Molarity original} \times \text{diluted factor}$$

~~d~~ ~~\*~~ 10

$$\cancel{\ast} \text{ diluted factor} = \frac{\text{original}}{\text{diluted}}$$

- a) gases behave ideally : ① low pressure  
 ② High Temp  
 ③ low d/wt

$$\rightarrow PV = nRT$$

$$\Rightarrow PV = \left( \frac{\text{Mass}}{\text{M.M.}} \right) \times R \times T$$

P: atm

T: K

V: L

M.M.: g/mole

Mass: g

torr  $\equiv$  mmHg

( $\approx 760$ )

atm

10)

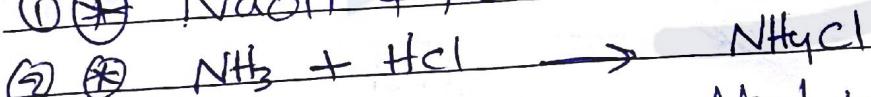
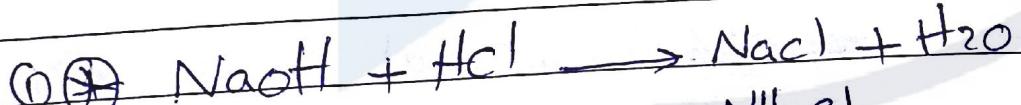
Heat lost = Heat gained

$$q_{\text{warm water}} = [q_{\text{cold water}} + q_{\text{cal}}]$$

for cold water

$$(m \times S \times \Delta t)_{\text{warm water}} = [(mS\Delta t)_{\text{cold}} + (C \times \Delta t)_{\text{cal}}]$$

cal constant



A)  $(P_{hot})Heat$  lost by hot water =  $m \times S \times \Delta t$

$$(q_{hot}) = \cancel{m} \times \cancel{S} \times \cancel{\Delta t} = \cancel{m} \times \cancel{S} \times \cancel{\Delta t}$$

Heat gained by cold water =  $(lost - cold)$

$$(C) \text{ constant} = \frac{\text{Heat gained by cold water}}{\Delta t \text{ for cold water}}$$

B)  $(q_{sol})Heat$  gained by solution =  $m \times S \times \Delta t$

$$(q_{sol}) = \cancel{m} \times \cancel{S} \times \cancel{\Delta t} = C \times \Delta t$$

Heat of the reaction =  $-(q_{sol} + q_{cold})$

$\Delta H$  = Heat of the reaction (kJ)

1 kJ of water  $\approx$  NaOH  $\approx$  HCl

$$* V + V = 2V$$

$$* m + m = 2m$$

c) same part (B)

\* to determine the third equation enthalpy



$$\Delta H_{\text{third}} = \Delta H_{\text{(first)}} + (- \Delta H_{\text{(second)}})$$

"Hess's Law"