

Aim: To estimate strength of the given oxalic acid in g/L by using approximately N/50 potassium permanganate as an intermediate solution.

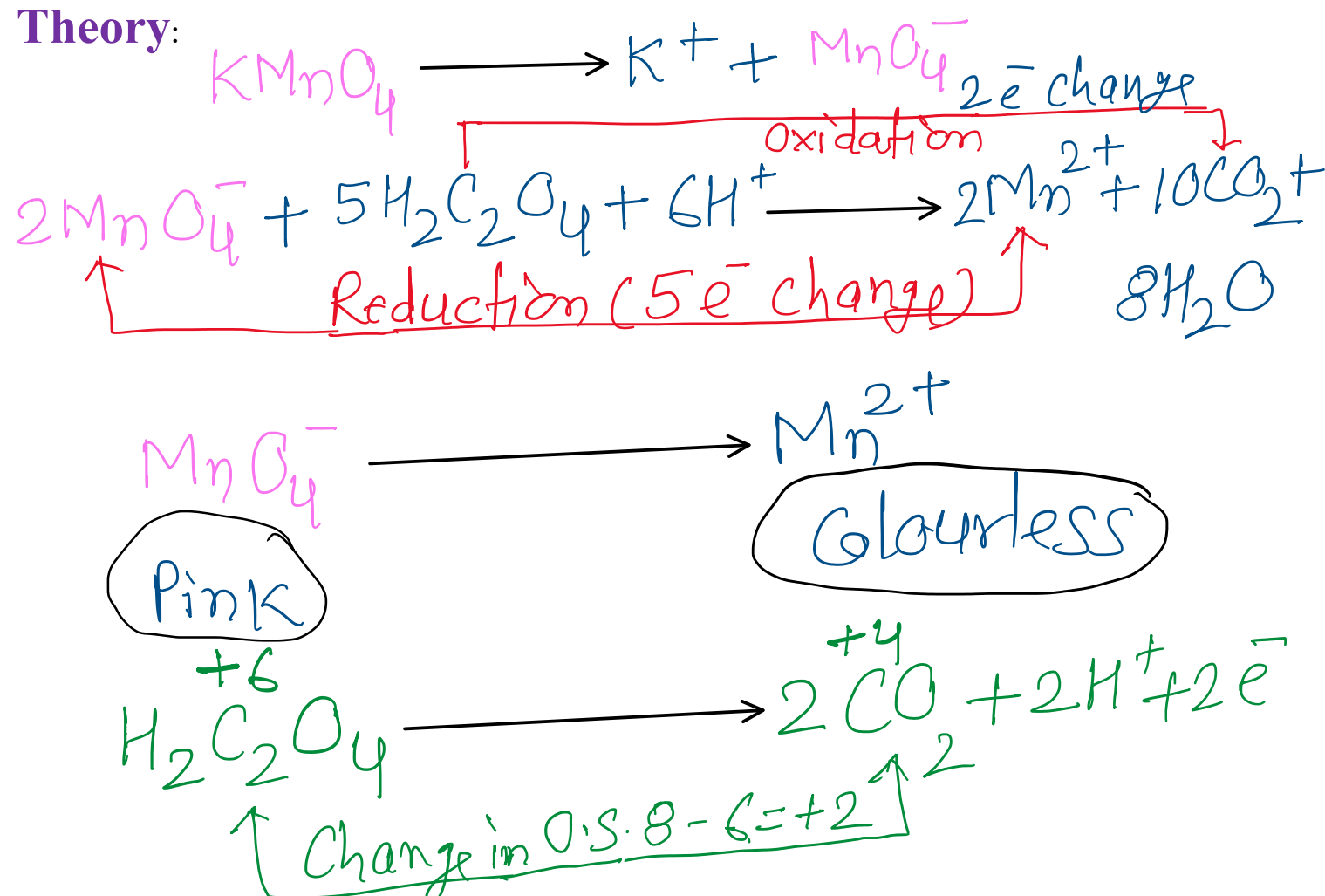
Sunday, 30 August 2020

8:17 PM

Requirements: Standard flask, conical flask, burette, pipette, burette stand, beaker, wash bottle, spatula etc.

Solutions and Reagents: Oxalic acid, Potassium permanganate and dilute sulphuric acid.

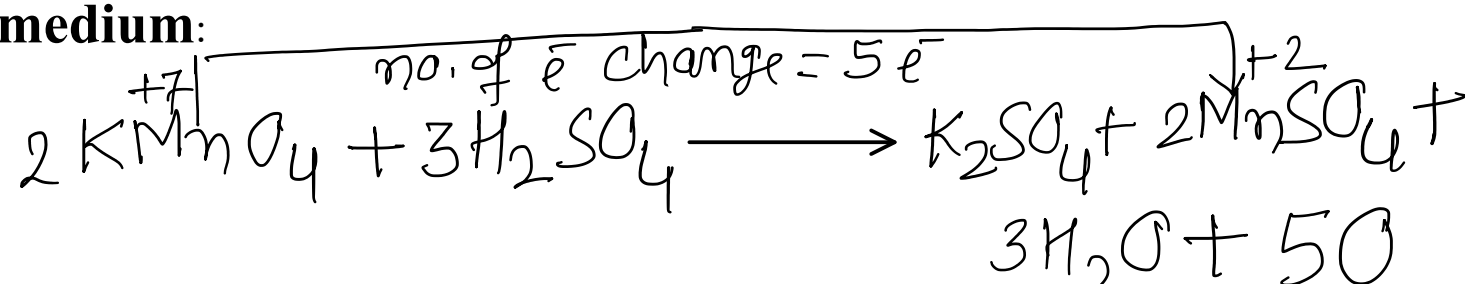
Theory:



Indicator: Potassium permanganate is a self indicator.

Colour Change at End Point: Colourless to pink

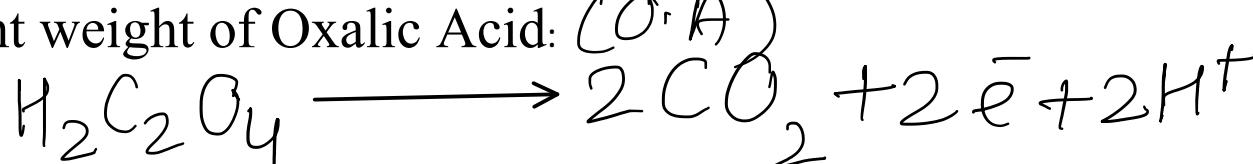
Equivalent Weight of Potassium Permanganate in acidic medium:



$$\begin{aligned} \text{Equivalent wt. of KMnO}_4 &= \frac{\text{Molecular wt.}}{5} \\ &= \frac{158}{5} \end{aligned}$$

$$\text{Equivalent wt. of KMnO}_4 = 31.6 \text{ in acidic medium}$$

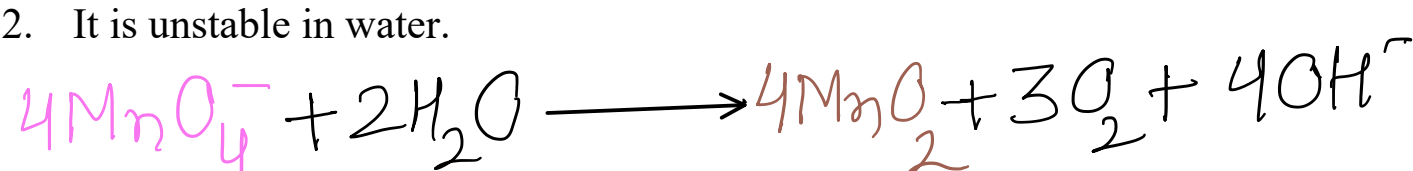
Equivalent weight of Oxalic Acid: (O.A.)



$$\text{Equivalent wt of O.A} = \frac{\text{Molecular wt.}}{2}$$

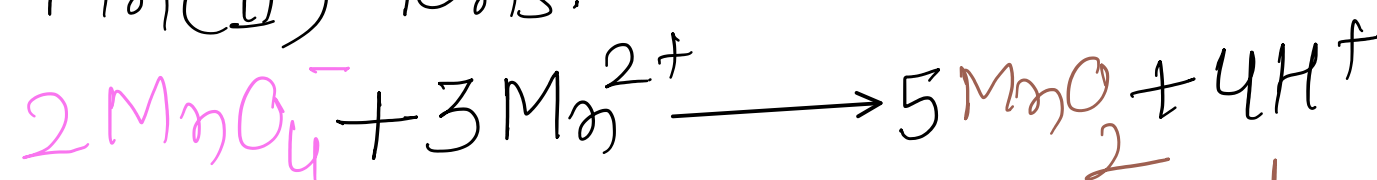
Why potassium permanganate is a secondary standard?

1. Potassium permanganate has some impurities of manganese dioxide.
2. It is unstable in water.



→ The presence of MnO_2 catalyses the auto-decomposition of MnO_4^-

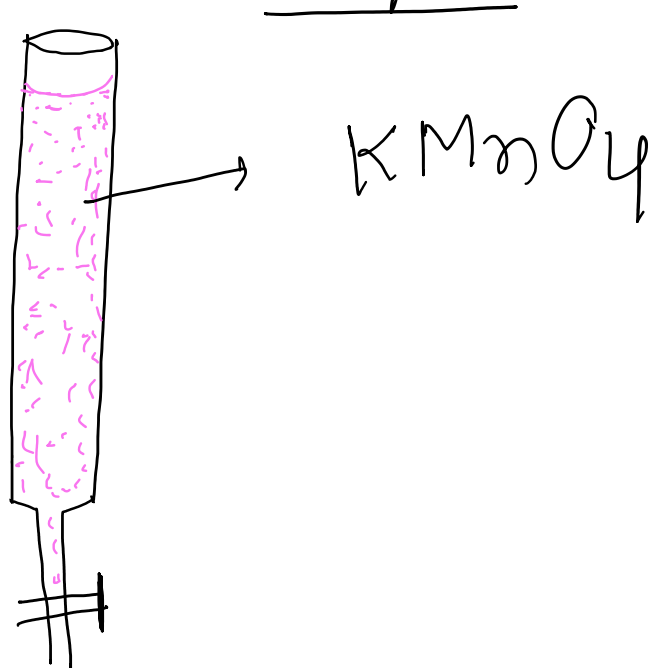
→ MnO_4^- is unstable in presence of Mn(II) ions.



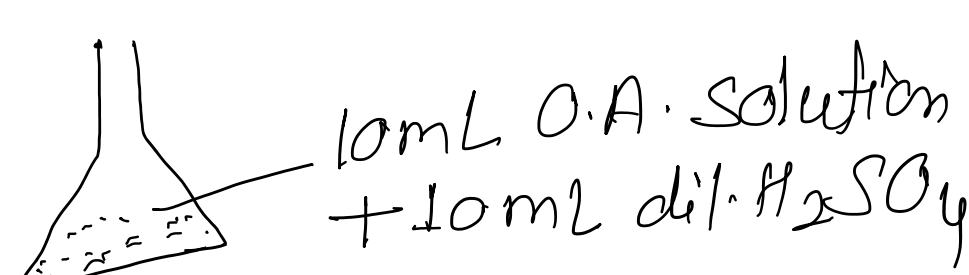
→ This reaction is slow in acidic medium, but it is very fast in neutral medium.

→ Because of these reasons KMnO_4 is always prepared with some amount of dilute acids.

Procedure: Step 1. fill the burette with KMnO_4 .

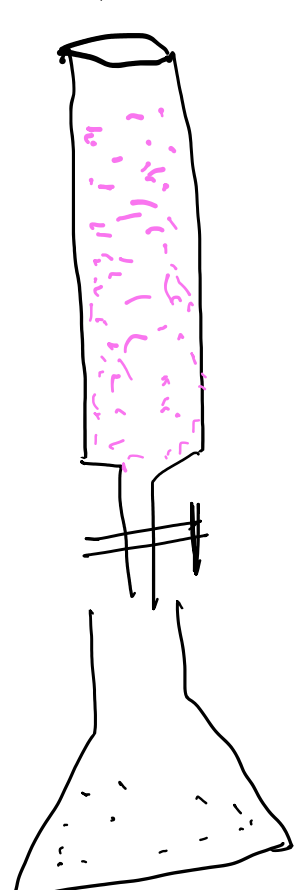


Step 2. Pipette out 10mL given oxalic acid solution in a clean conical flask and add to it 10mL dil. H_2SO_4 .



Step 3. Heat the contents gently to about 60°C .

Step 4. Titrate with KMnO_4 with constant stirring.



Step 5. At the end point faint pink appear which persists for 30s.

