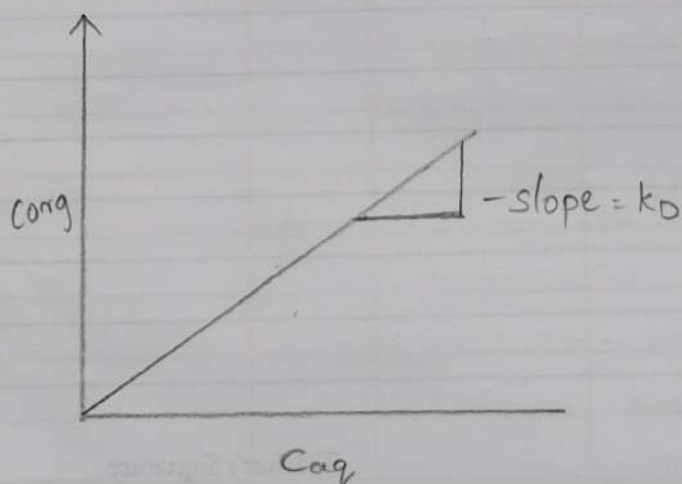


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S.No	Volume of M/20 hypo for org layer V_{org}	Volume of M/600 hypo for org layer	Corrg	C_{aq}	$K_D = \frac{C_{org}}{C_{aq}}$
1.	25.	12.5	0.258	0.024	10.74
2.	21.5	14.1	0.215	0.02	7.67
3.	17.8	10.5	0.178	0.04	8.47
4.	17.4	9.8	0.174	0.015	9.17
5.	13.5	9.2	0.135	0.018	7.5

Graph:- A Graph is plotted by taking C_{org} on y-axis & C_{aq} x-axis a straight line is passed through the origin is obtained. The slope of the straight line is equal to K_D (distribution co-efficient)



Distribution of I_2 between CCl_4 & H_2O

Aim:-

To determine the partition co-efficient of I_2 b/w CCl_4 and H_2O .

Apparatus:-

Reagent bottle, 2 burette & 1, 2 pipettes conical flask.

Chemicals:-

Saturated solution of I_2 in CCl_4 pure $m/20$ and $m/600$ hypo solution, starch solution.

Principle:-

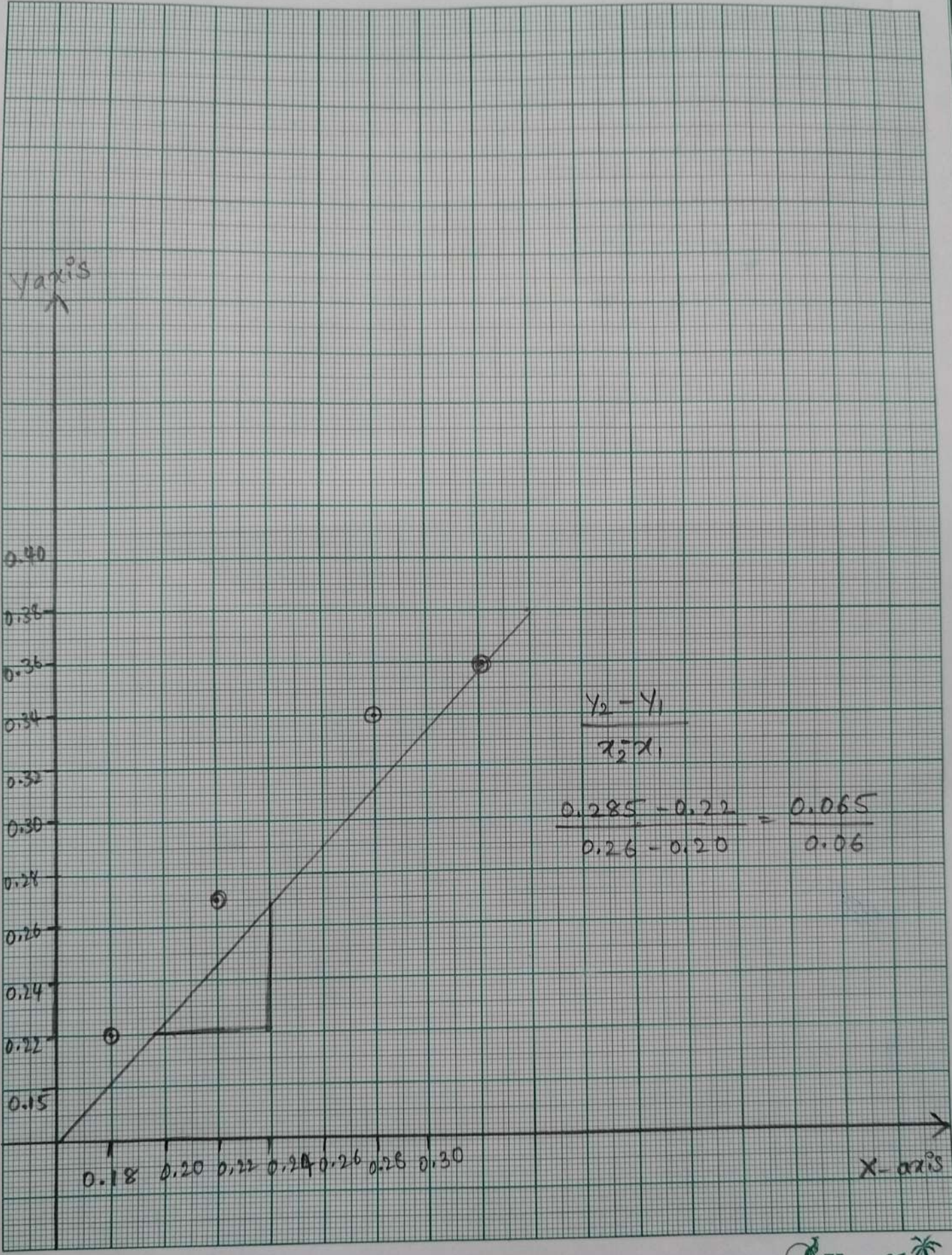
A common solute when added to a system of two immiscible liquids distributes itself in a definite concentration ratio. I_2 gets distributed itself between CCl_4 & H_2O separately. The partition co-efficient can be determined by tetra agents.

$K_D = \frac{\text{concentration of solute in organic phase} - \text{con of } I_2 \text{ in } CCl_4 \text{ layer}}{\text{con of solution in aqueous phase} - \text{con of } I_2 \text{ in } H_2O \text{ layer}}$

Procedure:-

Take 50 ml of I_2 in CCl_4 & 50 ml of H_2O in clean reagent bottle.

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$$\frac{y_2 - y_1}{x_2 - x_1}$$

$$\frac{0.285 - 0.22}{0.26 - 0.20} = \frac{0.065}{0.06}$$

* shake the bottle for about 10 minutes and allow it to settle for 5 minutes. The two layers, get separation the lower layer is CCl_4 layer & upper layer is aqueous layer.

* Get 10 ml aqueous layer & pipette out into a conical flask & titrated against $\text{M}/600$ hypo solution using starch as an indicator. The end point being the disappearance of blue color. The titra values noted as V_{aq} .

* 10 ml of organic layer (CCl_4) is pippered out into another conical flask & titrated against $\text{M}/50$ hypo solution. When the blue colour just disappear as & the titrated values is noted as V_{org} .

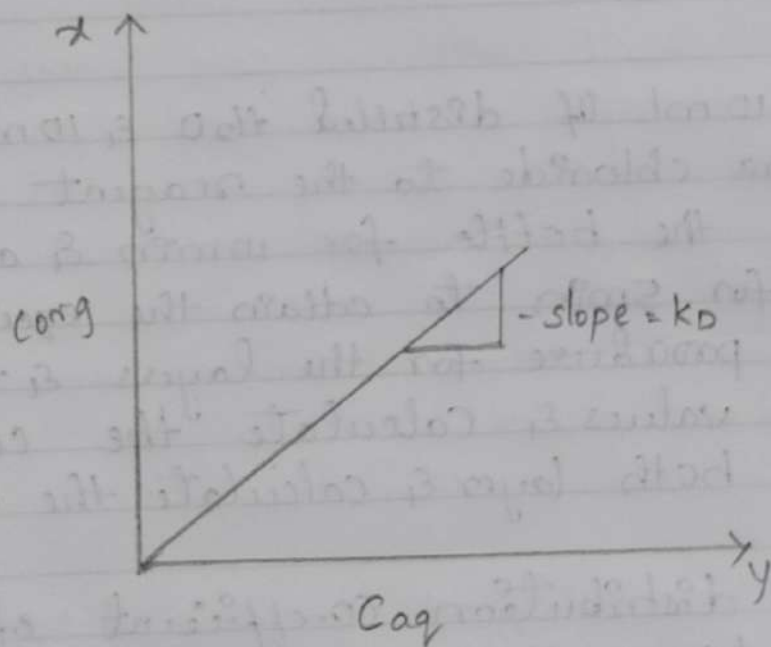
* Now add 10 ml of distilled H_2O & 10 ml of pure carbon tetra chloride to the reagent bottle and shake the bottle for 10 min & allow it to settle for 5 min to attain the equilibrium. Repeat the procedure for the layers & tabulate the titrate values & calculate the concentration of I_2 in both layer & calculate the results.

Result:-

The distribution co-efficient of I_2 b/w CCl_4 & H_2O .

S.NO	vol of NaOH for organic layer	vol of NaOH for aq layer	C_{org}	C_{aq}	$K_D = \frac{C_{org}}{C_{aq}}$
1.	43	36	0.43	0.36	1.19444
2.	37	30	0.37	0.30	1.2333
3.	34	24	0.34	0.24	1.416667
4.	26	23	0.26	0.26	1.150434

Graph:- A graph is plotted by taking C_{org} on x-axis & C_{aq} on y-axis a straight line is passed through the origin is obtained. The slope of the straight line is equal to K_D (distribution co-efficient)



Distribution of Acetic acid b/w n-butane & H₂O

Aim:-

To determine the distribution co-efficient of acetic acid b/w n-butanol & H₂O.

Chemicals:-

5% Acetic acid, 0.1 NaOH, n-butanol, phenol, phallen indicator.

Apparatus:-

Reagent bottle, conical flask, burette 10ml pipette

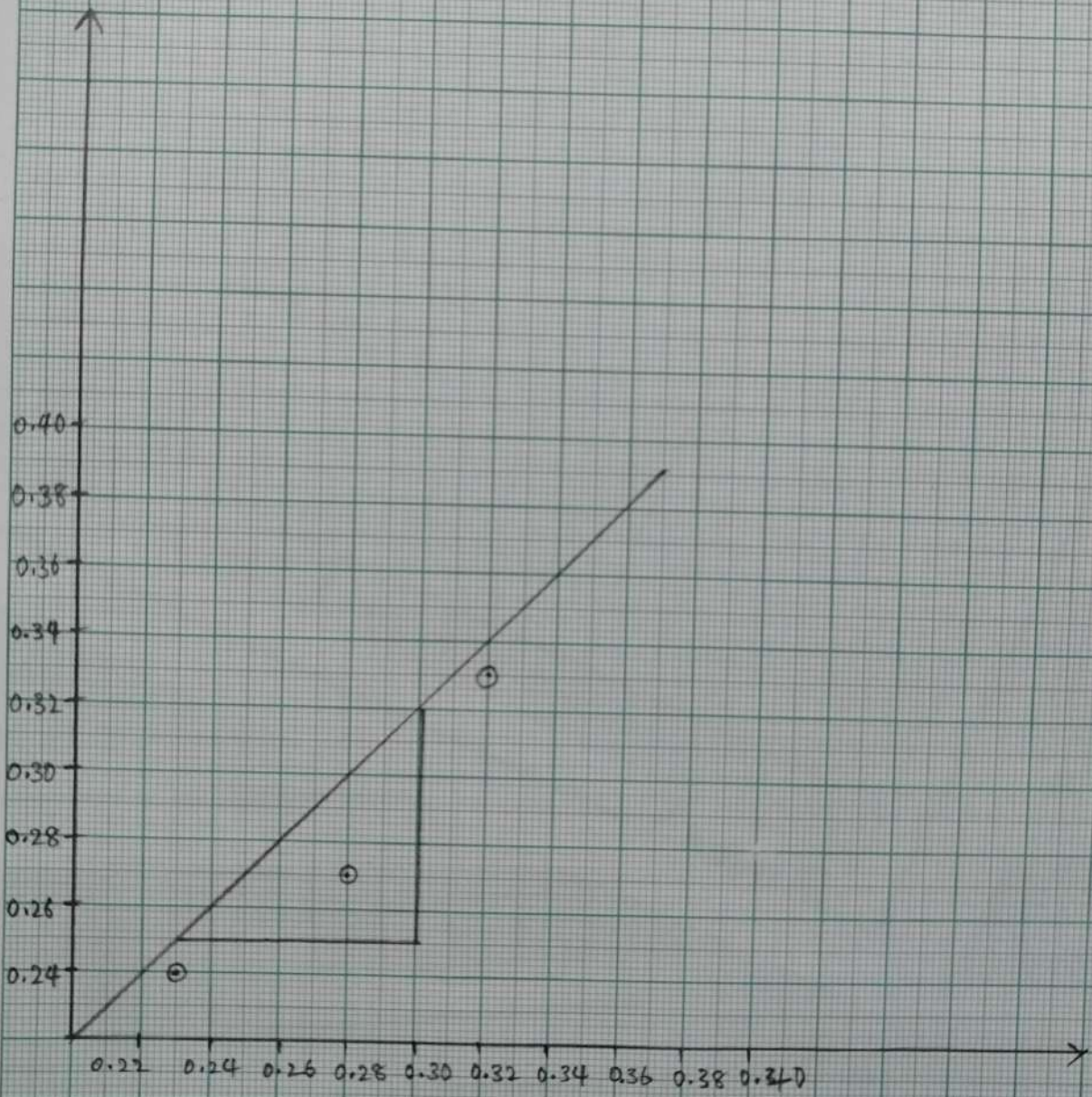
Principle:-

When a solid (or) liquid then the solute distribution two immiscible liquid & then the solute distributes itself in a such way that ratio of its constant temperature. If C_1, C_2 are then concentration & of solute in two solvents then the distribution position co-efficient K_D is given by.

$$K_D = \frac{\text{con. of solute in organic phase} = C_{org}}{\text{con. of solute in organic phase} = C_{org}}$$

Procedure:-

Take in a clean reagent bottle & take 50 ml of n-butanol & 50ml of 5% acetic acid.



- * cork the bottle & shake well without spilling of the solution for about 10 mins. Then open the cork & keep the bottle & acid and for 5 mins without disturbing, so that the two layers separate out.
- * Fill the burette with 0.1 M NaOH & clamp it to stand.
- * Pipette out 10 ml of organic layer & 10 ml of aqueous concial flask from the reagent bottle into two clean conical flask, add 2-3 drops of phenol pthalin indicator & titrate them against 0.1 M NaOH until the solution turns to pale pink in colour the titrated values are noted as V_{org} & V_{as} .
- * Now add the 10 ml of n-butanol & 10 ml of dist H_2O in the reagent bottle & shake well about 10 min keep a side 5 min without disturbing.
- * The procedure is repeated for three or more sets of readings & K_D is calculated.

Results:-

The distribution co-efficient of acetic acid b/n n-butanol & H_2O .

S. NO	Volume of NaOH (ml)	conductivity
1)	0	0.15
2)	1	0.45
3)	2	0.95
4)	3	1.35
5)	4	1.75
6)	5	1.98
7)	6	2.37
8)	7	2.63
9)	8	2.90
10)	9	3.03
11)	10	4.80
12)	11	5.31
13)	12	5.93
14)	13	6.54
15)	14	7.55
16)	15	6.54
17)	16	8.12
18)	17	8.48
19)	18	8.84

The procedure is repeated for three or more sets of readings & is calculated.

Results:
The titration is complete at 18 ml.

ELECTRO CHEMISTRY

Titration of weak acid against strong base
 CH_3COOH vs NaOH :-

Aim:-

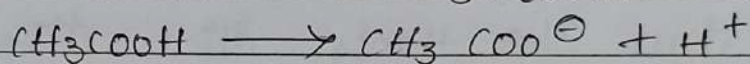
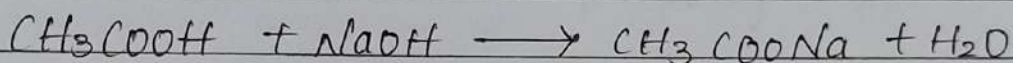
To determine the strength of a given weak acid (CH_3COOH) by conductometric titration with strong base (NaOH) of 0.01M.

Apparatus:-

Conductometer, burette, pipette, beaker & conical flask.

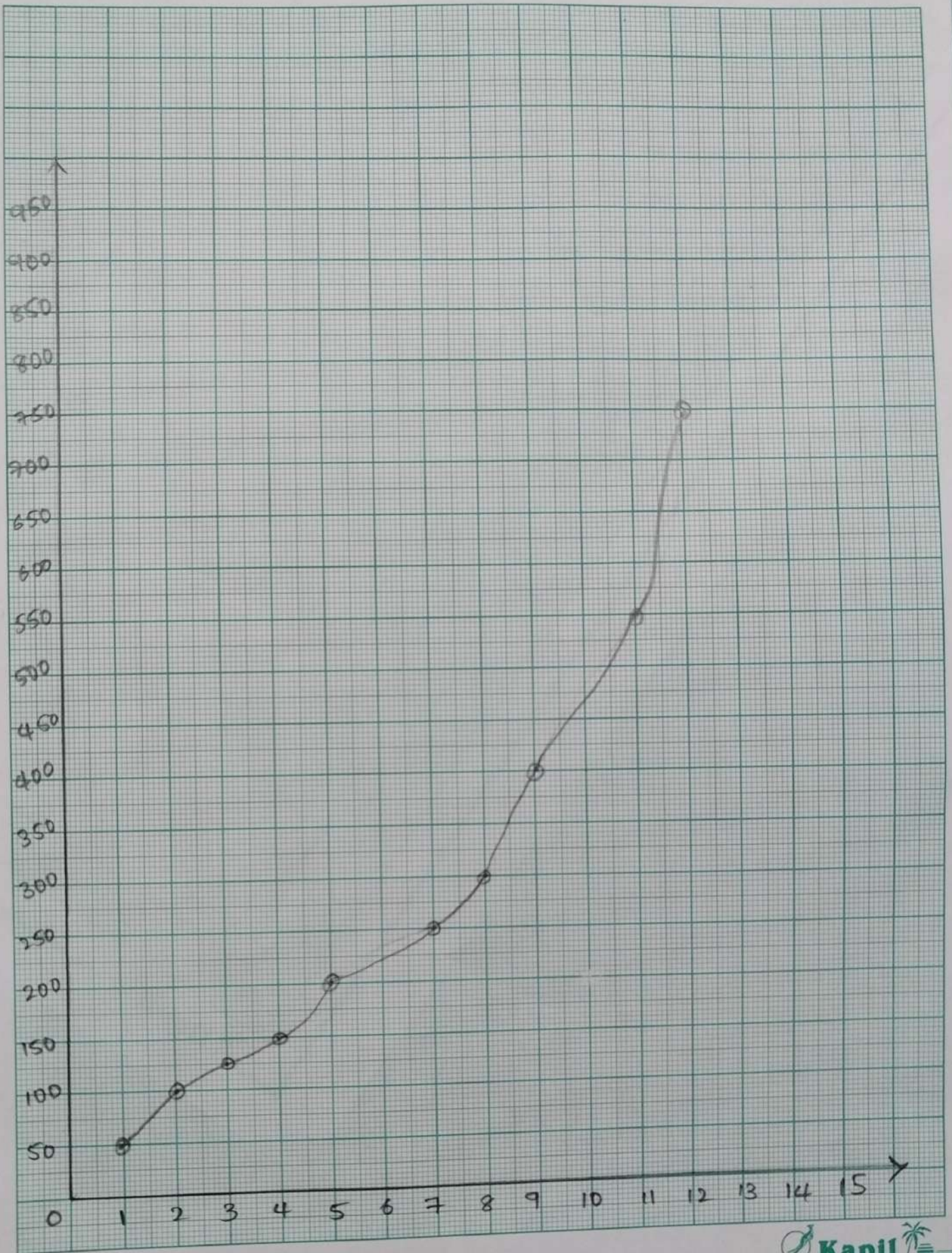
chemical flask:-

CH_3COOH solⁿ of NaOH solution

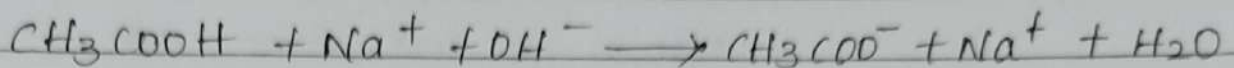


Principle:-

conductometric titration may be obtained as those titration in which end point is calculated (or) detected with the help of conductometer. Measure conductometer titration based on the fact of that conductances of solution at a constant temperature depends upon the number & molarity of ions present.



with strong base & initial conductance of acetic acid solution is low because of its poor dissociation on adding NaOH. solution highly minimized sodium acetate is formed.



Procedure:-

- Take 40 ml of given acid solution into a clean 100ml beaker deep the conductivity cell in the acid solution after proper cleaning with distilled H₂O if the cell is not properly dipped in acid solution & add little quantity of dist. H₂O so that the cell properly dipped.
- Fill up burette with NaOH solⁿ note the conductance of solⁿ before expect any addition of NaOH solution from the burette repeat the titration to get minimum 10 readings after the end point is reached.
- Tabulate the observed conductance & plot a graph by taking conductance on y-axis & volume of base on x-axis point of intersection of two required neutralise acid solution. Then concentration

of acid by using the following formula can be calculated.

$$M_1 V_1 = M_2 V_2$$

M_1 \longrightarrow con of CH_3COOH

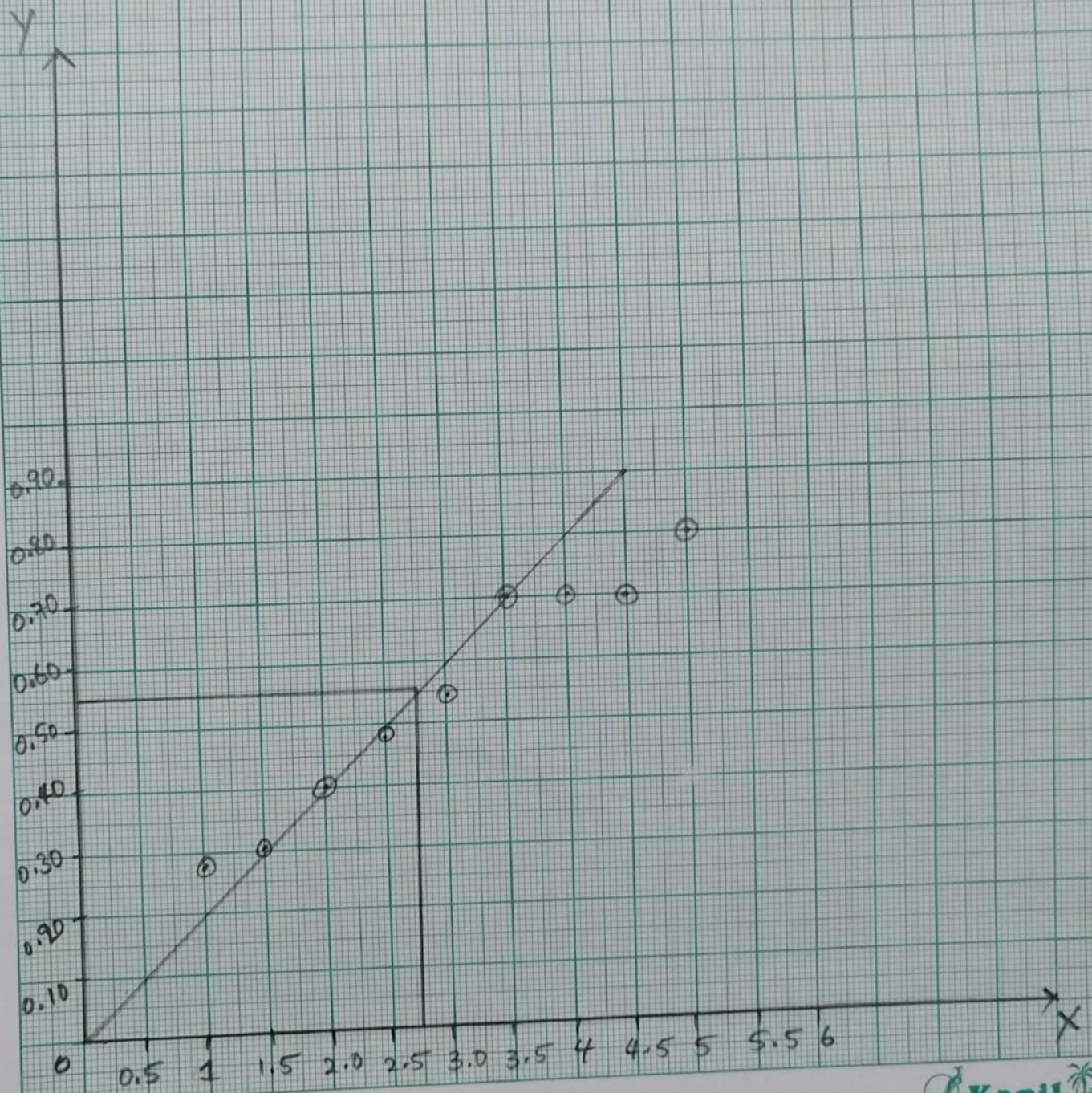
V_1 \longrightarrow vol of CH_3COOH

M_2 \longrightarrow con of NaOH

V_2 \longrightarrow vol of NaOH .

Result:—

The molarity of CH_3COOH obtained from the conductance titration is extracted is
0.1 M.



:- CALORIMETER :-

Verification of Beer's law for kmno_4 solⁿ

Aim:-

Verification of Beer's law by using kmno_4

Apparatus:-

Test tubes, cuvettes, calorimeter.

Chemicals:-

Potassium permanganate solution

Principle:-

According to Beer's law when monochromatic light is passed out through medium solution the rate of decrease in intensity of incident radiation with distance travelled is directly proportional to the initial intensity & concentration of substance absorbing light.

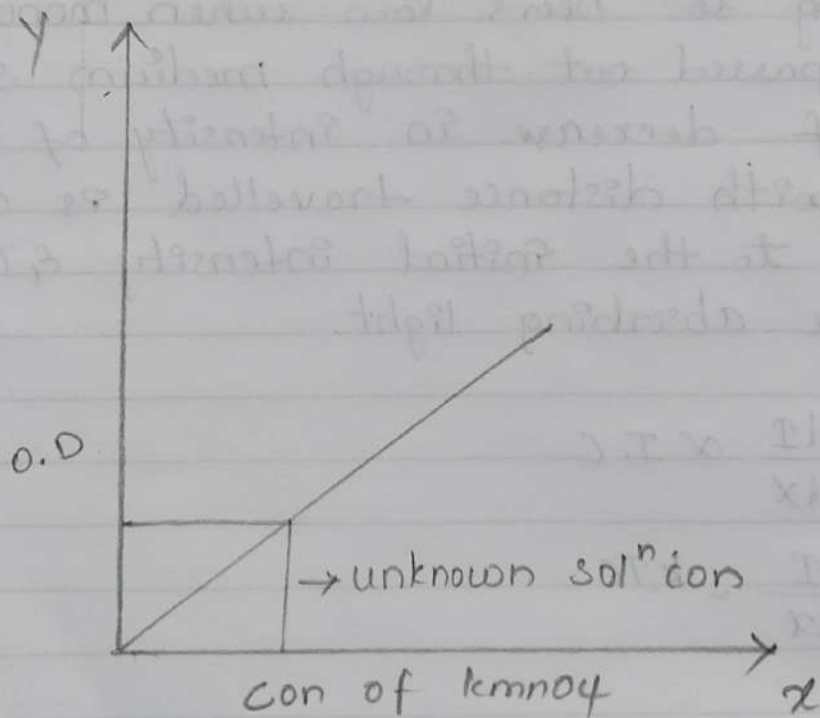
$$- \frac{dI}{dx} \propto I \cdot c$$

$$- \frac{dI}{dx} = k \cdot I \cdot c$$

$$- \frac{dI}{I} = k \cdot c \cdot dx$$

S.No	vol of kmno_4 in ml	vol of H_2O in ml	optical density	con of kmno_4
1,	10	0	0.80	5×10^{-4}
2,	9	1	0.70	4.5×10^{-4}
3,	8	2	0.68	4×10^{-4}
4,	7	3	0.67	3.5×10^{-4}
5,	6	4	0.54	3×10^{-4}

Graph - Graph is drawn by taking O.D on y-axis & con of kmno_4 on x-axis.



On integration

$$I + \frac{I}{I_0} = k \cdot c [x]_0^x$$

$$\log \frac{I_0}{I} = \frac{k}{2.303} [x]_0^x$$

$$\log \frac{I_0}{I} = \frac{k}{2.303} cx$$

$$A = \epsilon \cdot cx$$

Where,

A = Absorbance

E = molar extinction coefficient

c = concentration

x = path length

Procedure:-

prepare a solution of 0.1 M $KMnO_4$ solution calibrate the colorimeter by setting to 0% absorbance inserting a glass cuvette containing dist water.

- Now take the solution under study in the cuvette & the filter at which maximum absorbance is obtained is selected for further study.

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- Now set the colorimeter at selected filter and prepare the following concentration of KMnO4 solution.

Result:

The given unknown KMnO4 solution concentration is 2.5%.

Adsorption :-

Adsorption of Acetic acid animal charcoal.

Aim:-

To verify the adsorption of Isotherms for acetic acid adsorption on animal charcoal.

Apparatus:-

Reagent bottle-3, Burette funnel, watching glass, filter paper, 10 pipette, conical flask.

Chemicals:-

0.1 acetic acid, 0.7m NaOH charcoal & 1gm of phenolphthalein.

Principle:-

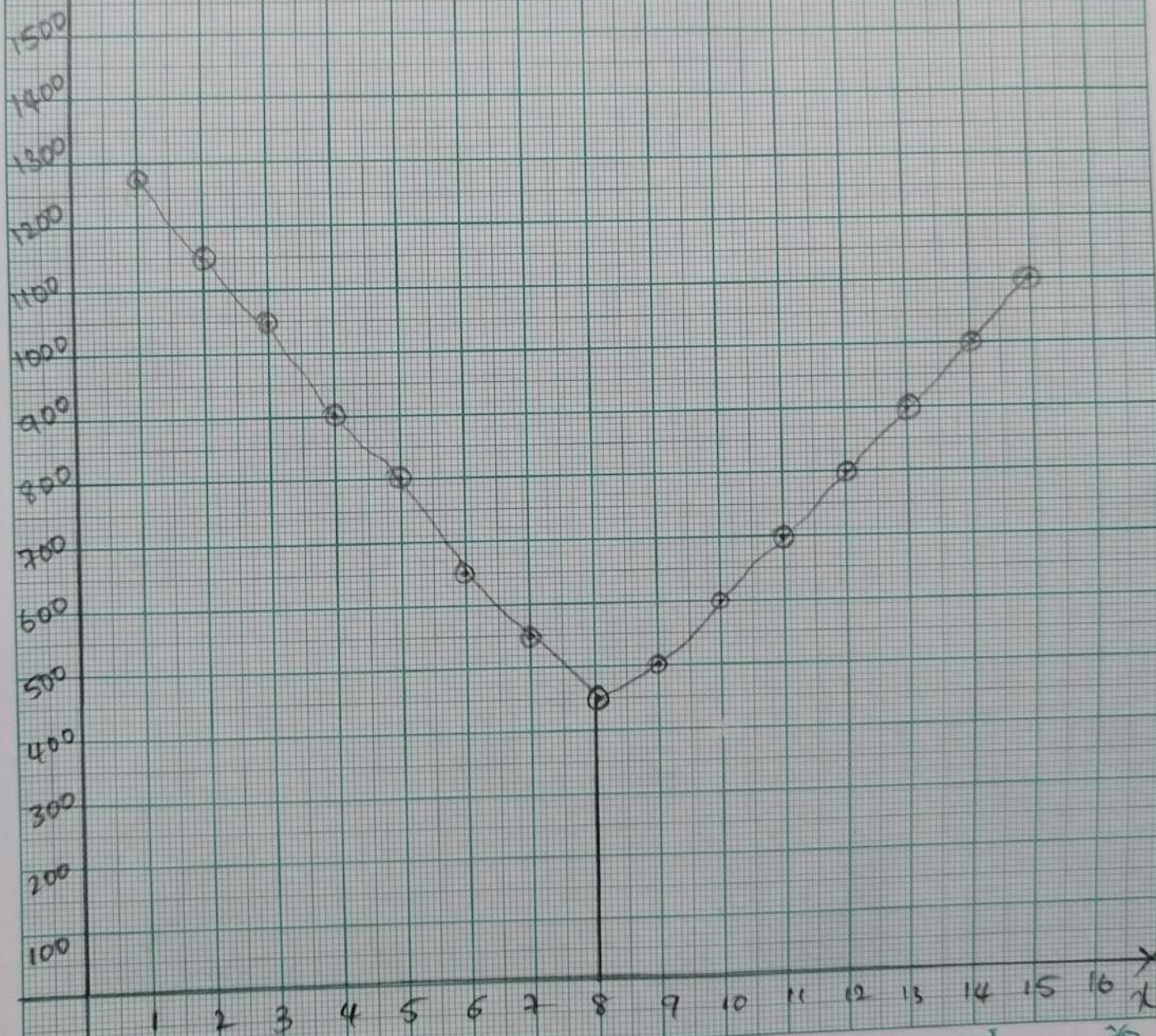
Freundlich proposed a relation b/w the amount of a solute adsorption on a definite amount adsorbent & the equilibrium conc. of the adsorbate in the solution according to which.

$$x/m = k \cdot c_e^{1/n} \rightarrow \log(x/m) = \log k + \frac{1}{n} \log c_e$$

where,

n \rightarrow amount of solute absorbed
 c_e \rightarrow equilibrium conc of adsorbent
 k \rightarrow constant.

Scale
x-axis - term - ml
y-axis - term - 100 ml



$n \rightarrow$ no. of layer acetic acid absorbed to the surface layer of charcoal.

Procedure:-

Take a clean reagent number than 1-5 prepare the following a solution mixture.

Fill the burette with 0.1 M NaOH solution. Pipette out 10 ml of stock solution 0.1 M acetic acid solution into a conical flask and add (or) 2 drops of phenolphthaleine indicator solution the end point is noted when a (length point) light pink colour is observed.

After 1 hour filter the content of each bottle separately through a filter paper number 4 which following reject the first 5 ml of filtrate and calculate & pipette out 10 ml of filtrate in to a clean conical flask.

And titrate it again with 0.1 M NaOH by adding 1 (or) 2 drops reagent repeat the procedure with the filtrate of other bottle and calculate the result.

Graph:-

A graph is plotted taking $\log(x/m)$ on y-axis & $\log(u)$ on x-axis is a straight line. The slope of the straight line is equal to γ -axis & the intercept is equal to $\log(c)$.

Result:-

The adsorption of isotherm is verified.

PHYSICAL CONSTANT

Determination of physical constant Surface tension.

Aim:-

To determine the surface tension of their given liquid using pylonometer.

Apparatus:-

pylonometer, rubber, tube, weight box.

Chemicals:-

Benzene, Carbon, tetrachloride, n-hexane

Principle:-

The surface tension of a given liquid can be calculated by using the equation

$$f = \frac{h_2}{n_1} = \frac{n_1 \times d_2}{h_2 \times d_1}$$

f_2 = The surface tension of a given liquid

h_2 = The no. of drops for the given liquid

d_2 = velocity of the given liquid.

n_1 = no. of drops of water

d_1 = The density of water

S_1 = The surface tension of water.

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S.NO	liquid	no. of drops n_1	no. of drops n_2	mean $\left(\frac{n_1+n_2}{2}\right)$	substance - tension	literate value	γ -error
1							
2							
3							
4							
5							
6							
7							
8							
9							
10							

The surface tension of a given liquid can be calculated using the equation

$$\gamma = \frac{m}{n} \times \frac{1}{2} \times \frac{1}{r}$$

1. The surface tension of a given liquid is the force per unit length acting along the surface of the liquid.

2. The surface tension of water is 72 dyne/cm.

Procedure:-

Salgonometer consists of a fine capillary tube with the center the upper & 10 mm ends of the bulb consists two marks say A and B.

- clean the salgonometer thoroughly with distilled H_2O followed by chromic acid and again with distilled H_2O followed by thoroughly the upper end slightly above the mark A and light the rubber tube with a pinch of clamp the salgonometer.
- Now discharge H_2O & dry the tube by using air current on in the given the repeat the process with other given liquids & tabulate the result.
- calculate the surface tension using the above given formula.

Results:-

Surface tension of given liquid
temperature $0^\circ C$ $d \text{ gm/cm}$.

PHYSICAL CONSTANT

Determination of constant velocity

Aim:—

To determine the velocity of given liquids using ostwald viscometer

Apparatus:—

ostwald, viscometer, lamp, breaker

Chemicals:—

Benzene, carbon tetrachloride, n-benzene

Theory:—

Postulate derived the fundamental equation to measure the co-efficient of velocity as below

$$b = \frac{\pi r^4}{8 \eta l} \frac{P}{\Delta \rho}$$

η = co. efficient of velocity

V = volume of the liquid.

l = length of narrow tubes radius of tube

t = The temperature under the above the equation can be reduced as

$$\frac{h_2}{h_1} = \frac{\rho_1 t_1}{\rho_2 t_2}$$

Since the hydrostatic pressure is directly proportional to the density of liquid.

$$\frac{h_2}{h_1} = \frac{d_1 t_1}{d_2 t_2}$$

Procedure:—

- The Ostwald viscometer consists of a capillary tube a bulb "B" at its upper. There are two marks X & Y on the tube & above and below the bulb "A".
- Clean the viscometer thoroughly with dist. H₂O followed by chromic acid and again with dist. H₂O.
- Set the viscometer & introduce measured volume of given organic liquid into the large bulb "B" now suck the liquid up into the bulb A through the capacity a rubber tube attached to upto a level begin the capillary plate, then stop cork when the liquid reaches upto mark X & note temperature when it reaches mark Y response

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the same to get concordant reading.

- Once the density given complete of organic compound liquid & water co-efficient of velocity of water then value for the liquid can be calculated.

Result:-

Co-efficient of velocity of the given compound is at temperature 0°C .