

CONDUCTOMETRY

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CONDUCTOMETRY

- INTRODUCTION
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- CONDUCTOMETRIC TITRATIONS
- ADVANTAGES & DISADVANTAGES OF CONDUCTOMETRIC TITRATIONS
- APPLICATIONS OF CONDUCTOMETRY

● INTRODUCTION:

➤ It is an electrochemical method of analysis concerned with electrical conductance through an electrolyte solution .

(or)

➤ It is defined as determination or measurement of the electrical conductance of an electrolyte solution by means of a conductometer .

❖ electric conductivity of an electrolyte solution depends on :

1. Type of ions (cations, anions, singly or doubly charged)
2. Concentration of ions
3. Temperature
4. Mobility of ions

- PRINCIPLE:

- Based on the conductance of electrical current through electrolyte solutions similar to metallic conductors
- The electric conductance in accordance with ohms law which states that the strength of current(i) passing through conductor is directly proportional to potential difference & inversely to resistance.

$$i = V/R$$

- Important definitions & relations
 - Conductance
 - Specific conductance
 - Molar conductance
 - Equivalent conductance
 - Resistance
 - Specific resistance

- Conductance:(G)

- ❖ ease with which current flows per unit area of conductor per unit potential applied & is reciprocal to resistance(R)

$$G = I/R$$

- Specific conductance (K):

- ❖ conductance of the body of uniform length(l) & uniform area cross section(A)

$$K = 1/R \times 1/A$$

- Molar conductance: (\wedge)

- ❖ Conductance of a solution containing 1 mole of the solute in 1000 cm of the solution which placed between two parallel electrodes which are 1 cm apart

$$\wedge = 1000/C$$

- Equivalent conductance: (\wedge_{eq})

- ❖ specific conductance of the solution containing 1gm equivalent of solute in 1000cm³ of solution.

$$\wedge_{eq} = 1000k / c_{eq}$$

- Resistance (r):

- ❖ Is a measure of the conductors opposition to the flow of electric charge

$$R = 1/G$$

- Specific resistance:(ρ)

- ❖ Is resistance offered by a conductor of unit length and having unit cross section

$$R \propto l/A$$

• Instrumentation

The instrument used for measurement of conductance are known as conductometers

It consists of :

1. Current source
 - Alternating current source
2. Conductivity cells
 - Wide mouthed cells
 - Cell for reactions producing precipitates
 - Dip type cells
3. Electrodes

CURRENT SOURCE:-

1. Mechanical high frequency AC generator by Washburn .
2. Vreeland oscillator by Taylor and Acree.
3. Vacuum tube oscillator by Hall & Adams.
 - ❖ When electrical potential is applied across electrodes two processes occur.
 - Ions accumulate near the electrodes.
 - Transfer of charge through the interface.
 - Note : DC current is not employed in conductance measurement because
 1. Electrodes become polarised leading to high cell resistance.

● Conductivity cells:-

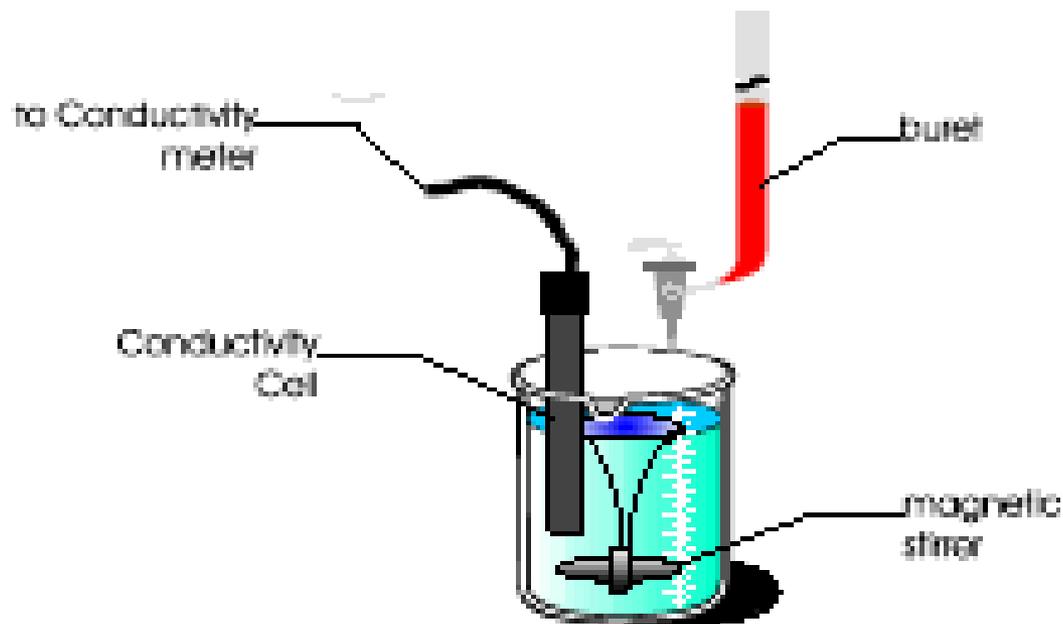
- Made of pyrex or quartz and are fitted with two platinum electrodes.
- Should be placed in vessel containing water to maintain constant temperature
- Types :
 1. Wide mouthed cell
 2. Cell for reactions producing precipitation
 3. Dip type cells



- Wide mouthed cell:-

- Measurement of low conductance
- Wide mouthed fitted with an ebonite cover which has provisions for platinum electrodes and burettes

- Cell for reactions producing ppts:
 - Mainly used for ppt reactions
 - Also wide mouthed fitted with ebonite cover which has provisions for burette ,electrode , stirrer
 - Stirrer may be mechanical or magnetic



- Electrodes:

- Platinum sheets, each of 1 cm^2 are fixed at distance of 1 cm
- The surface is coated with platinum black to avoid polarization effects and increase effective surface area.
- Platinisation of electrodes is done by coating solution of 3% chlorplatinic acid and lead acetate on it to get uniform coating
- Electrodes usage depends on conductivity and concentration
- If conc is low then electrodes should be largely and closely packed

- Measurement:-

- The instrument used to measure conductance is called conductance bridge or conductometer
- Classical circuit employed for measurement is wheatstone bridge
- All other work on this principle
- Various types are:
 1. Kohlrausch conductance bridge
 2. Direct reading conductance bridge
 3. Phillips conductance bridge
 4. Mullard's conductance bridge
 5. Pye's conductance bridge

- Kohlrausch conductance bridge:
 - Consists of a meter bridge XY with fixed resistors r' & r'' at both ends. One arm of bridge consists of resistance box 'R' & other arm with conductivity cell 'C'. Detector D is head phone while inductance coil 'J' is AC source which is operated by battery.
- Direct reading conductance bridge:-
 - In this head phone is replaced by magic eye which is electronic device

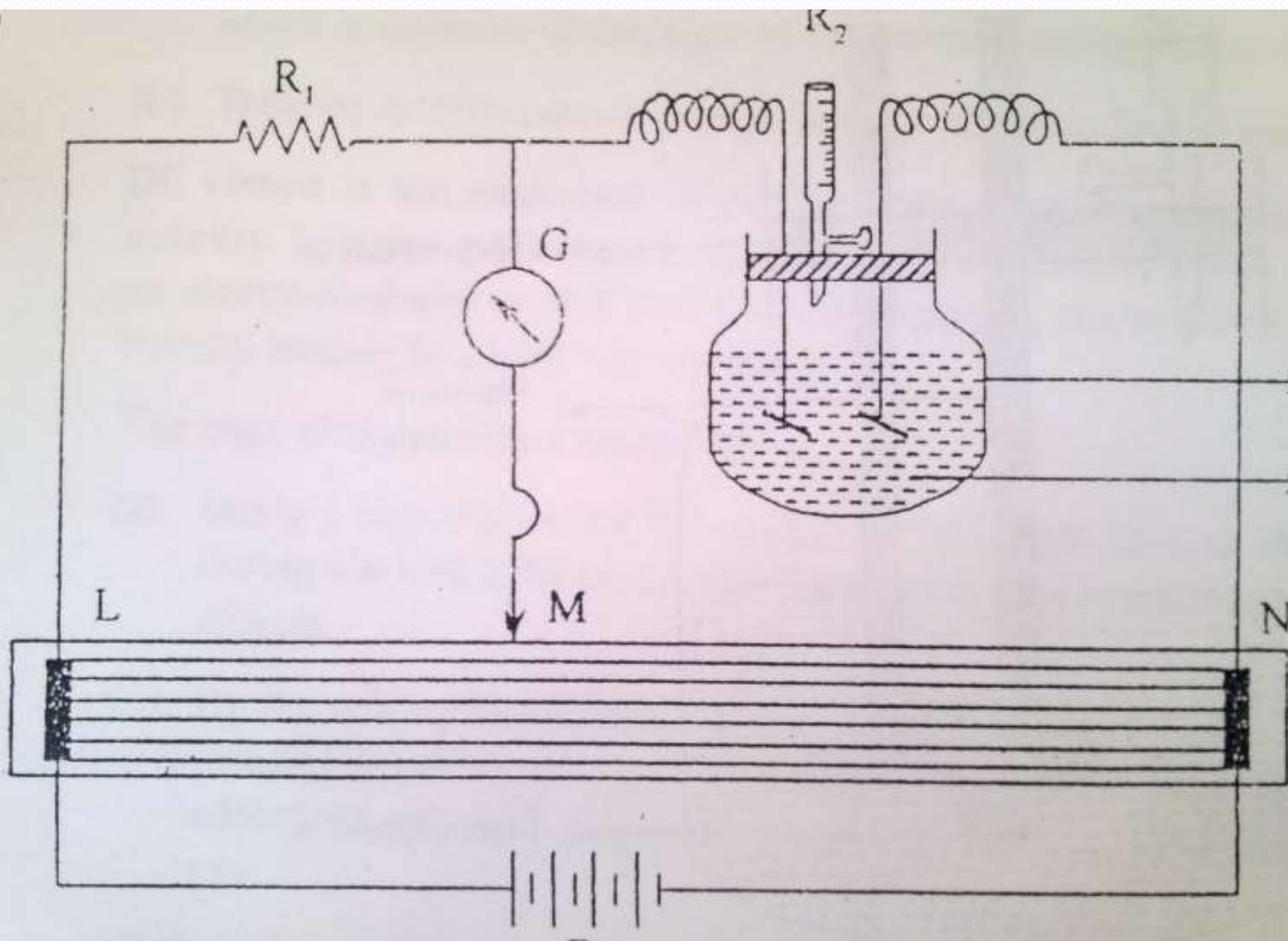
- The set up for Measurement :

consists of meter bridge LN attached to standard resistance R_1 & unknown resistance R_2

cell is connected to standard resistance to one side , meter bridge LN at other. The sliding contact with galvanometer (G) can be moved on the wire of meter bridge by means of jockey (M) so that resistance of unknown is balanced with that of standard. When galvanometer shows null deflection, the resistance of unknown is measured by following equation:

$$ML / NL = R_2 / R_1$$

$$R_2 = ML / NL \times R_1$$



Conductivity cell

Solution

Meter bridge

- Hence conductivity of unknown solution:

$$1/R_2 = NL/ML \times R_1$$

The measured conductivity ($1/R_1$) is not always equal to the specific conductivity of solution, because the physical configuration of platinum electrode i.e, length and area of electrodes varies from one another. Hence conductivity of solution is obtained by calculating a factor called “cell constant”.

- Cell constant:

Defined as ratio of distance between the two electrodes(l) to the area of electrodes(A)

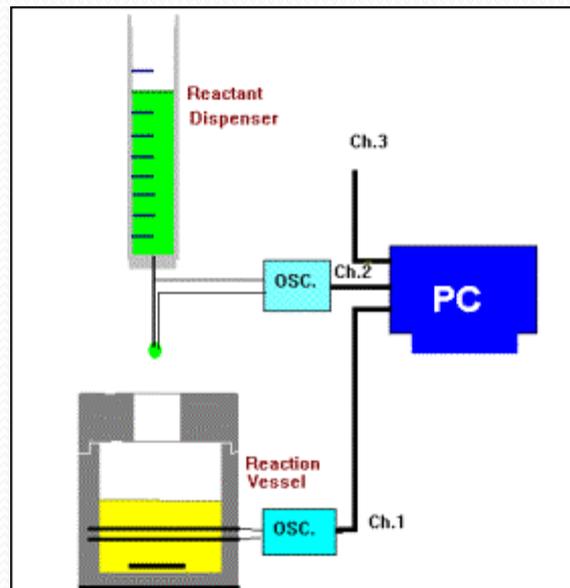
There fore,

$$\theta = l / A$$

- CONDUCTOMETRIC TITRATIONS:

- INTRODUCTION:

- Is process of qualitative chemical analysis in which conc of sample is determined. Which is done by adding areagent(titrant) of known conc in measured volumes to the sample (anylate)



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- TYRES OF CONDUCTOMETRIC TITRATIONS:
 - Acid –base or neutral titrations
 - Replacement or displacement titrations
 - Redox titrations
 - Precipitation titrations
 - Complexometric titrations
 - Non-aqueous titrations

1. ACID- BASE OR NEUTRAL TITRATIONS:

➤ STRONG ACID-STRONG BASE

- EG: HCL vs NaOH

➤ STRONG ACID-WEAK BASE

- EG: HCL vs NH_4OH

➤ WEAK ACID-STRONG BASE

- EG: CH_3COOH vs NaOH

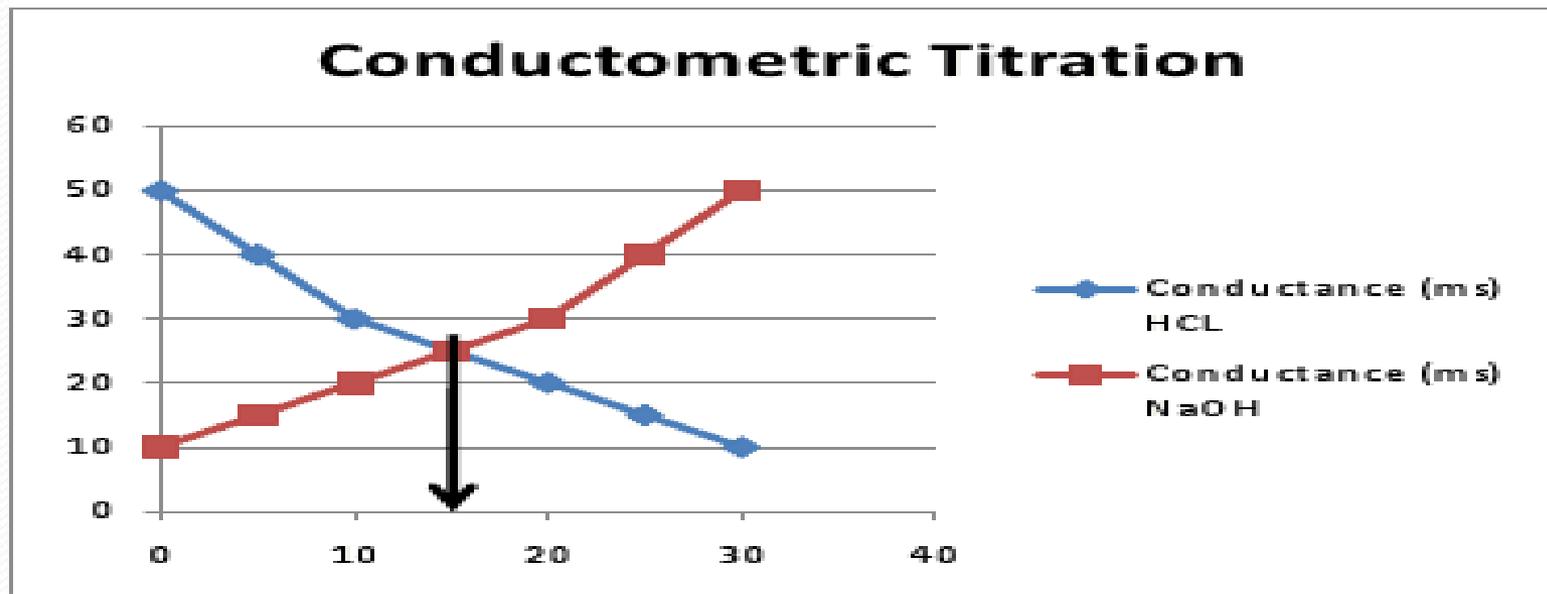
➤ WEAK ACID -WEAK BASE

- EG: CH_3COOH vs NH_4OH

- Strong acid strong base:

- Fall in conductance due to replacement of high conductivity Hydrogen ions by poor conductivity sodium ions
- Rise in conductance due to increase in hydroxyl ions

Strong Acid-Strong Base

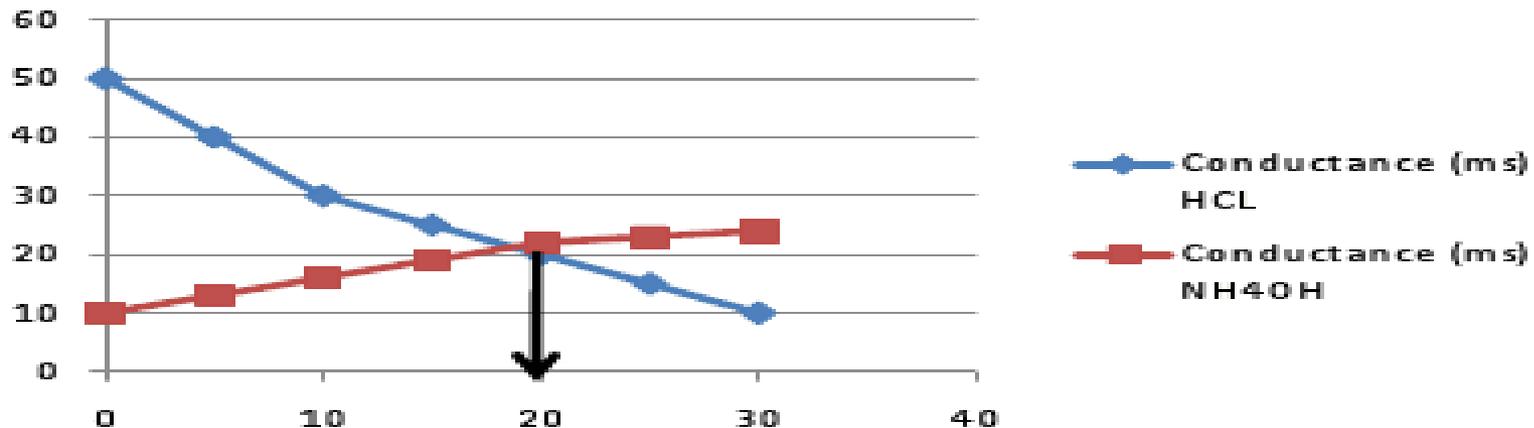


- Strong acid- weak base:

- Fall in conductance due to replacement of hydrogen by ammonium ions
- Conductance remain constant due to supression of NH_4OH by NH_4CL

Strong Acid-Weak Base

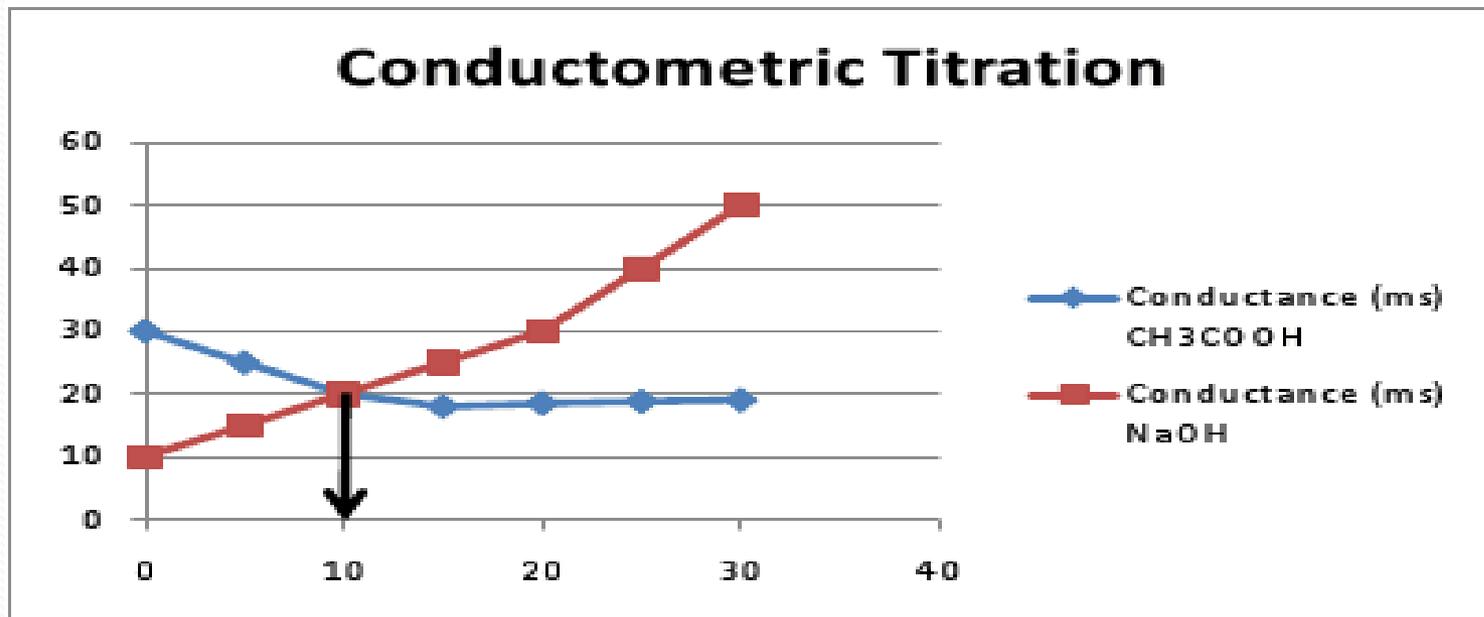
Conductometric Titration



- Weak acid –Strong base:

- Initial decrease in conductance followed by increase due to NaOH
- Steep rise due to excess of NaOH

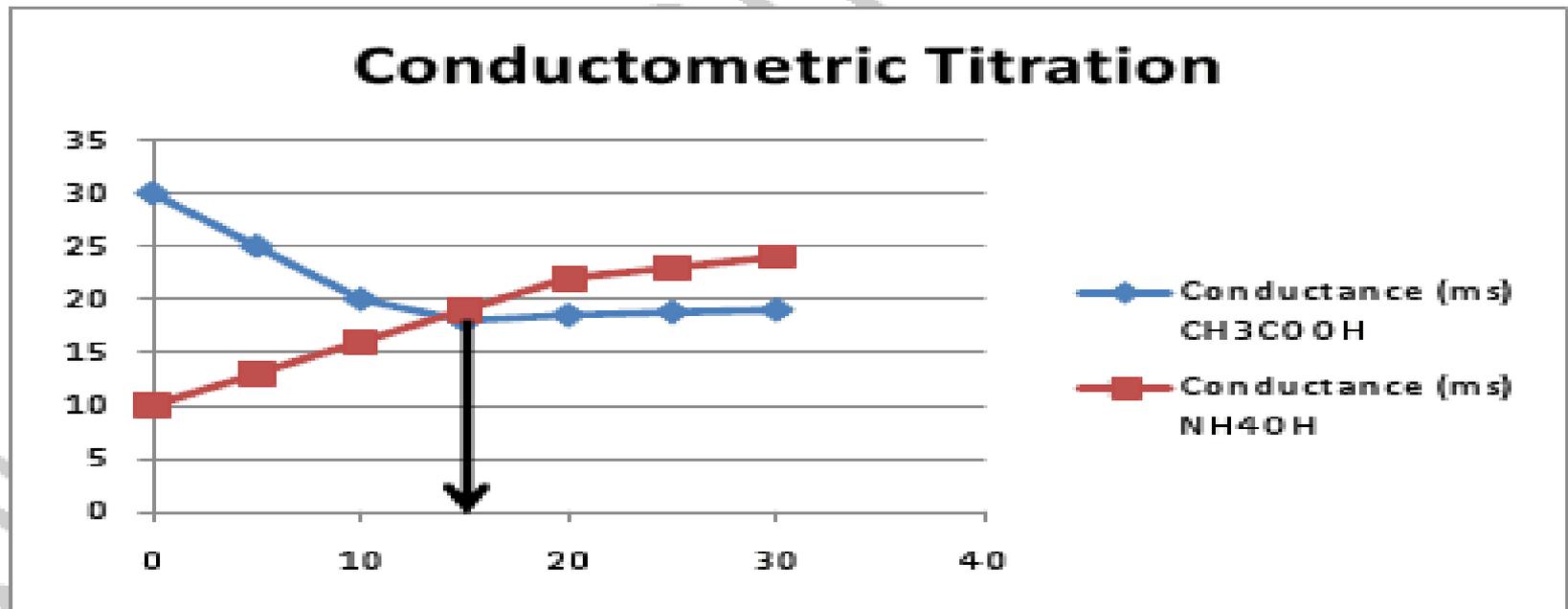
Weak Acid-Strong Base



- Weak acid- weak base:

- Increase in conductance due to excess of CH_3COOH
- Constant conductance due to supression of NH_4OH by CH_3COOH

Weak Acid-Weak Base



- **ADVANTAGE OF CONDUCTOMETRIC TITRATIONS:**
 1. Does not require indicators since change in conductance is measured by conductometer
 2. Suitable for coloured solutions
 3. Since end point is determined by graphical means accurate results are obtained with minimum error
 4. Used for analysis of turbid suspensions, weak acids, weak bases, mix of weak & strong acids

- Disadvantages of conductometric titration:

1. Increased level of salts in solution masks the conductivity changes, in such cases it does not give accurate results
2. Application of conductometric titrations to redox systems is limited because, high concentrations of hydronium ions in the solution tends to mask the changes in conductance

- Applications :

1. Check water pollution in rivers and lakes
2. Alkalinity of fresh water
3. Salinity of sea water (oceanography)
4. Deuterium ion concentration in water- deuterium mixture
5. Food microbiology- for tracing micro organisms
6. Tracing antibiotics
7. Estimate ash content in sugar juices
8. Purity of distilled and de - ionised water can determined
9. Solubility of sparingly soluble salts like AgCl , BaSO_4 can be detected
10. Determination of atmospheric SO_2 , estimation of vanillin in vanilla flavour

References

- *Instrumental analysis* by A. Skoog, F. James Holler and Stanly R. Crouch.
- *Text book of pharmaceutical analysis*, third edition by Dr.S.Ravi sankar.



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THE END

