

CY 104_Applied Chemistry

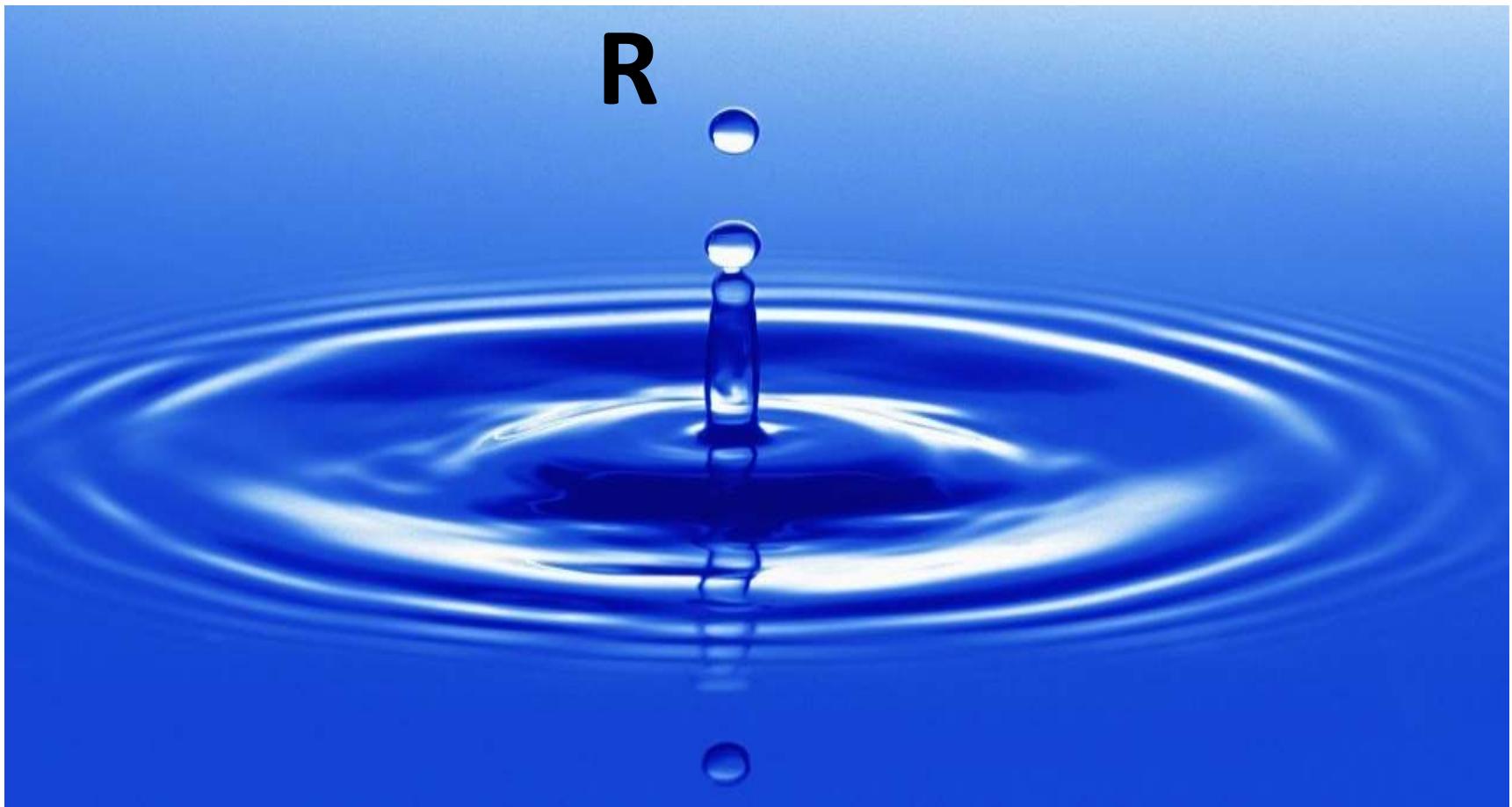
by

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WATE

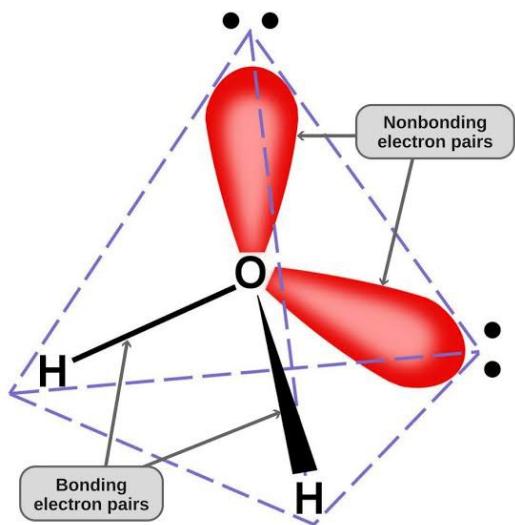
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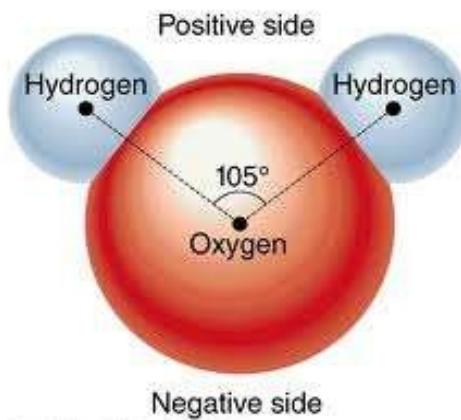
- Water is essential for life. It covers 2/3 of the earth's surface and every living thing depends upon it.
- The human body is comprised of over 70% water, and it is a major component of many body fluids including blood, urine, and saliva.



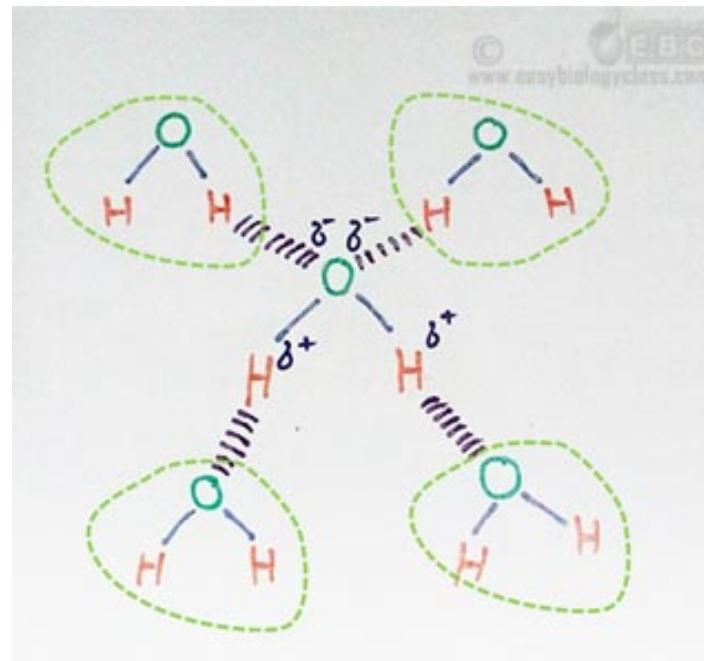
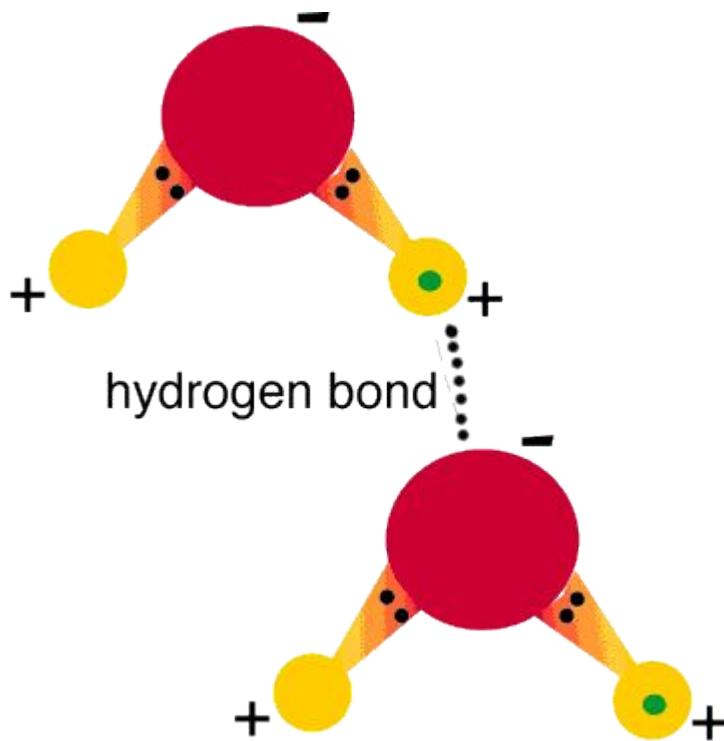
- Water is formed when two hydrogen atoms bond to one oxygen atom.
- With a total atomic weight of 18 daltons
- The structure of the electrons surrounding water is tetrahedral, resembling a pyramid.
- The angle between H-O-H bond is 104.45 degree.



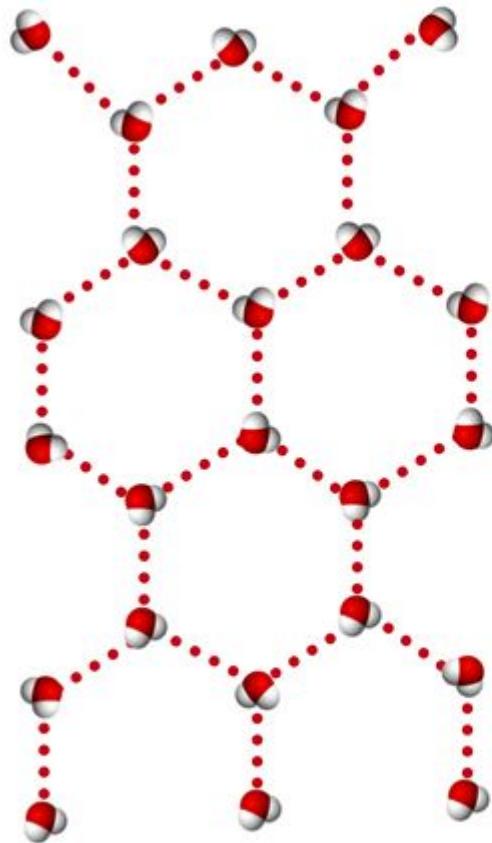
- There is a great concentration of electrons around the nucleus of the oxygen than around the hydrogen.
- Therefore, the hydrogen end is slightly positive and the oxygen end is slightly negative.



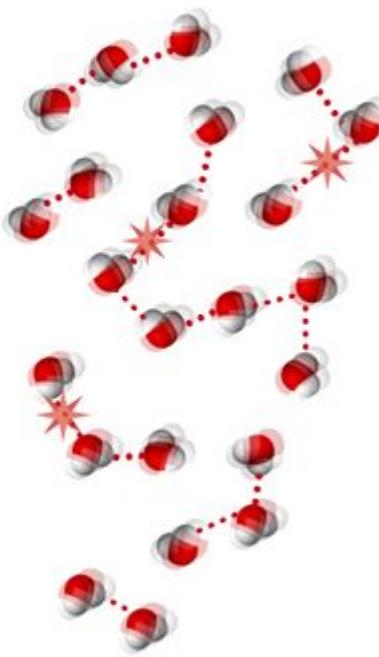
- Hydrogen bond exists between a highly **Electronegative atom** (like oxygen in another water) of a polar molecule and a **hydrogen**
- Weak bond, but strong in great numbers.**



Single water molecule can form FOUR hydrogen bonds at a time in Liquid water



a) Solid water (ice)



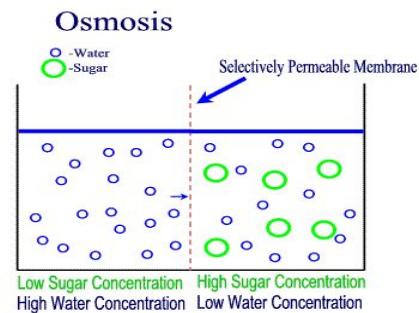
b) Liquid water

Properties of water

Physical Properties

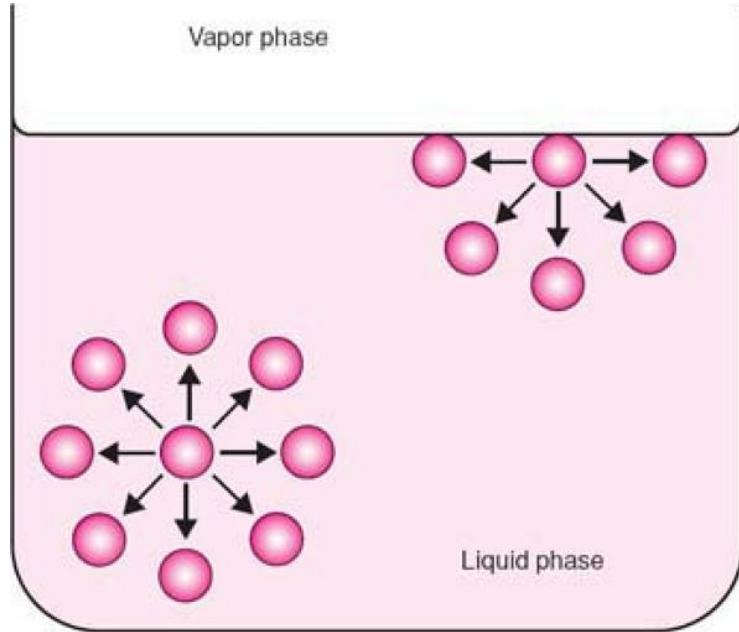
- ✓ Polar molecule: Water is a polar molecule because of the way the atoms bind in the molecule with more positive(+) charges on one side of the molecule and more negative(-) charges on the other side of the molecule. In other words the hydrogen atoms group on one side of the molecule making that more positive, such that there are more electrons from the oxygen atom on the other side of the molecule.
- ✓ Solubility: Water dissolves salts such as NaCl by hydrating and stabilizing the Na and Cl ions, weakening the electrostatic interactions between them. The same factors apply to charged biomolecules, Water readily dissolves such compounds by replacing solute-solute hydrogen bonds with solute-water hydrogen bonds, thus screening the electrostatic interactions between solute molecules.

- ✓ Osmosis: Osmosis is the passage of water from a region of high water concentration through a semi-permeable membrane to a region of low water concentration. Osmosis, transfer of a liquid solvent through a semipermeable membrane that does not allow dissolved solids (solutes) to pass.

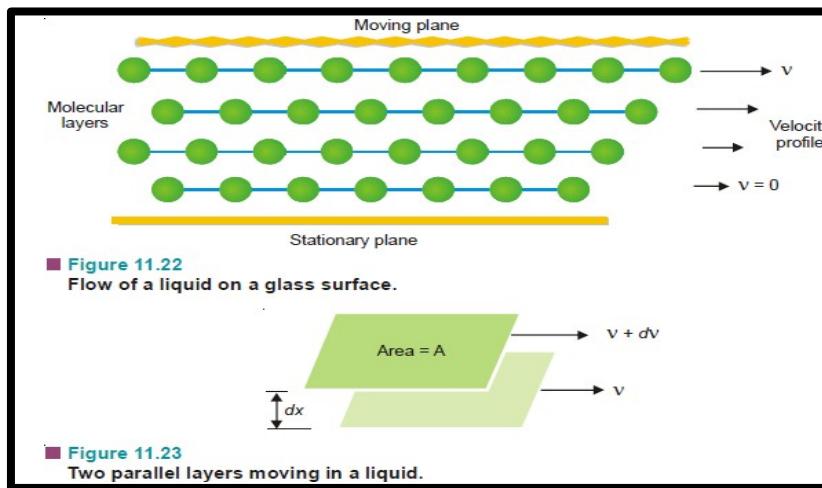


- ✓ Surface tension : Molecules at the surface (i.e., at the liquid–air interface) can only develop attractive cohesive forces with other liquid molecules that are situated below and adjacent to them. They can develop adhesive forces of attraction with the molecules constituting the other phase involved in the interface, although, in the case of the liquid–gas interface, this adhesive force of attraction is small. The net effect is that the molecules at the surface of the liquid experience an inward force toward the bulk. Such a force pulls the molecules of the interface together and, as a result, contracts the surface, resulting in a surface tension.

72 dynes/cm



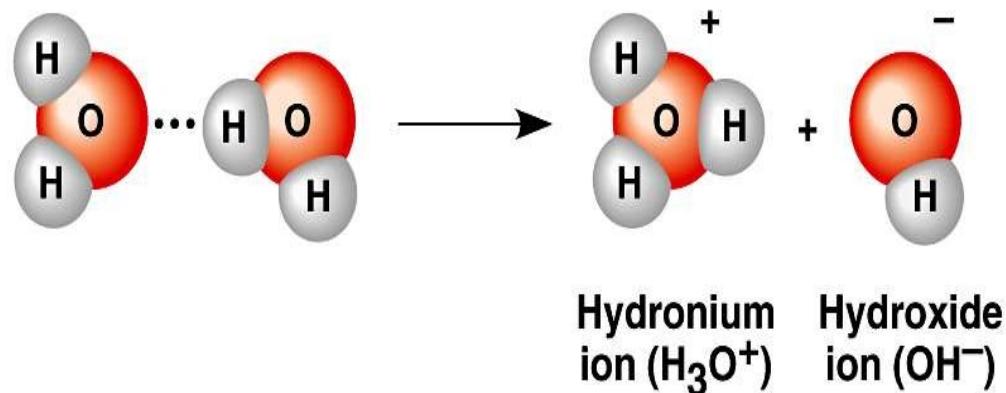
- ✓ Viscosity: When a shearing force is applied to a liquid, it flows. However, the forces of friction between the layers offer resistance to this flow.



Water has a **viscosity** of 0.0091 poise at 25 °C

Chemical properties

- ◎ Occasionally, a hydrogen atom shared by two water molecules shifts from one molecule to the other.
 - > The hydrogen atom leaves its electron behind and is transferred as a single proton - a **hydrogen ion** (H^+).
 - > The water molecule that lost a proton is now a **hydroxide ion** (OH^-).
 - > The water molecule with the extra proton is a hydronium ion (H_3O^+).



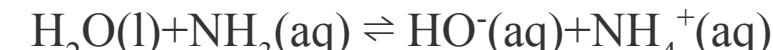
Water reacts with a lot of substances to form different compounds. Some significant reactions are as follows:

- **Amphoteric nature:**

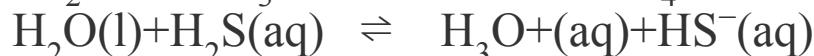
Water can act as both acid and base, which means that it is amphoteric in nature.

Example:

Acidic Behaviour:

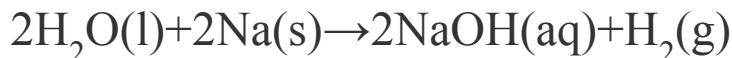


Basic Behavior:



- **Redox reactions:**

Electropositive elements reduce water to hydrogen molecule. Thus, water is a great source of hydrogen. Let us see an example in this case:



During the process of photosynthesis, water is oxidized to O_2 . As water can be oxidized and reduced, it is very useful in redox reactions.

- **Hydrolysis reaction**

Water has a very strong hydrating tendency due to its dielectric constant. It dissolves many ionic compounds. Some covalent and ionic compounds can be hydrolyzed in water.

Sources of

The Main Sources Of Water Are: **Water:**

- 1) Rain water
- 2) River water
- 3) Spring or well water
- 4) Sea water

1). **Rain water:** Rain water is the purest form of natural water. However, it dissolves considerable amount of gases (CO_2 , SO_2 , NO , NO_2 etc) and suspended solid particles from atmosphere, during its journey through it and becomes polluted.

2). **River water:** Rivers are formed by rain and spring waters. During its flow over the surface of land, it dissolves minerals of the soil such as chlorides, sulfates, bicarbonates of sodium, calcium, magnesium ions etc..

3) **Spring or well water or Lake water:** it contains constant chemical composition.

The minerals present in the lake water in the form dissolved form and high quantity of organic matter.

4) **Sea water:** It is the most impure form of natural water. It contains larger percentage of the dissolved salts (above 3.5%) out of which about 2.6% is NaCl. The NaCl which is present in the dissolved form in sea water will come out as NaCl crystals due to evaporation of sea water. The other salts present in the sea water are sulphates of sodium, bicarbonates of potassium, magnesium, calcium, bromides of potassium, magnesium etc.

5) **Underground water:** Spring & well waters are the underground water sources. They are in general clearer in appearance due to the filtering action of the soil.

They contain more of the dissolved salts generally, underground water is of high organic purity.

Types of water

1. Raw water:

- Untreated, unfiltered water
- Unsafe for human consumption
- Without treatment raw water can be used for farming, construction or cleaning purposes.

2. Cooling water:

- Cooling water system are widely used in industry to reject waste heat.
- Cooling water should be free from solid impurities which could damage pumps and cause blockages.

3. Boiler water:

- Water within a boiler, associated with piping, pumps and other equipments, that is intended for evaporation into steam.

4. Nuclear water:

- Water is used in processing (mining, milling, fuel fabrication) of fuel (Uranium).
- Nuclear cooling systems are designed so that if pipes begin to leak, local water runs into the plant rather than radioactive water leaking out.
- After being removed from the reactor the nuclear fuel is still very hot and requires storage both to cool down and to control risks of radiation poisoning.

Types of impurities in water:

The impurities present in water are classified as:

1). Dissolved impurities: Dissolved impurities may organic or inorganic. Inorganic impurities: the carbonates, bicarbonates, sulphates, chlorides of calcium, magnesium, iron potassium and aluminium.

Organic impurities: Organic water products, amino acids, proteins, etc.

Gases: O_2 , CO_2 , Oxides of nitrogen and sulphur, H_2S etc.

2) Suspended impurities: It is of two types:

1. Inorganic - sand & clay;
2. Organic – vegetable and animal matter.

3) Biological Impurities: Micro-Organisms like Pathogenic bacteria, fungi, algae etc

HARDNESS OF WATER

Hard Water : Those water which does not produce lather (or) very little lather with soap is called Hard Water

Soft Water : Soft water readily produce a lot of lather when mixed with little soap.

The Hardness of water is caused by the presence of dissolved salts such as Bicarbonates, Sulphates, Chlorides and Nitrates of bivalent metal ions like Ca^{+2} & Mg^{+2}

Soap is sodium/potassium salt of higher fatty acids like stearic, oleic and palmetic acids.

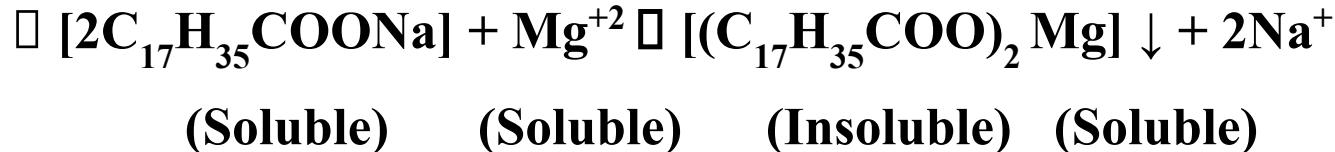
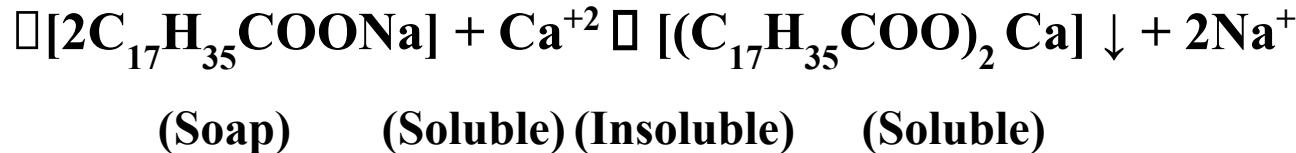
When soap is mixed with soft water lather is produced due to stearic acid and sodium stearate



\square Stearic Acid + Na-Stearate \square Formation of lather.

When soap comes in contact with HARD WATER,

Sodium stearate will react with dissolved calcium and magnesium salts and produce calcium stearate or magnesium stearate which is white precipitate



DISADVANTAGES OF HARDWATER / CAUSES OF HARDNESS:

The following are the disadvantages when hard water is used for various purpose:

(i) DOMESTIC USE:

- (a) Washing and Bathing : Hard water does not form lather easily with soap is wasted
- (b) Drinking : Hard water causes bad effects on our digestive system. Sometimes, stone formation takes place in kidneys
- (c) Cooking : The boiling point of water is increased due to the presence of salts.
Hence, more fuel and time are required for cooking.

(ii) INDUSTRIAL USE:

- (a) Textile Industry : Hard water causes wastage of soap. Precipitates of calcium and magnesium soap adhere to the fabrics and cause problem
- (b) Paper Industry : Calcium and Magnesium salts in water may effect the quality of paper.
- (c) Sugar Industry : Water containing sulphates, carbonates, nitrates affects the crystallisation of sugar.
- (d) Pharmaceutical Industry : Hard water may form some undesirable products while preparation of pharmaceutical products.

(iii) STEAM GENERATION IN BOILERS: For steam generation, boilers are employed. If hard water is used in boilers, It may lead to the following troubles

- (a) Boiler Corrosion
- (b) Scale and Sludge formation.
- (c) Priming and Foaming
- (d) Caustic embrittlement

TYPES OF HARDNESS

The hardness of water is of two types

- (1) Temporary hardness (or) Carbonate hardness
- (2) Permanent hardness (or) Non-Carbonate hardness
- (3) Temporary Hardness: Temporary hardness is caused by two dissolved bicarbonate salts $\text{Ca}(\text{HCO}_3)_2$ and $\text{Mg}(\text{HCO}_3)_2$.
The hardness is called “Temporary Hardness”
Because it can be removed easily by means of boiling.



(2) PERMANENT HARDNESS:

Permanent hardness of water is due to the dissolved chlorides, sulphates and nitrates of calcium and magnesium.

These salts are CaCl_2 , CaSO_4 , $\text{Ca}(\text{NO}_3)_2$, MgCl_2 , MgSO_4 , $\text{Mg}(\text{NO}_3)_2$

These Hardness cannot be removed easily by boiling. Hence it is called “Permanent Hardness”. Only chemical treatment can remove this hardness.

Total Hardness Of Water = Temporary Hardness + Permanent Hardness

DEGREE OF HARDNESS:

- The Concentration of hardness as well as non-hardness constituting ions are, usually expressed in the term of “Equivalent amount of CaCO_3 ”
- Since this mode permits the multiplication and division concentration, when required. The choice of CaCO_3 in particular is due to its molecular weight (m.wt) is “100” (Equivalent wt = 50), and moreover, It is insoluble salt that can be precipitated in water treatment.

- Therefore 100 parts by weight of CaCO_3 hardness must be equivalent to

1 162 parts by weight of $\text{Ca}(\text{HCO}_3)_2$ hardness

2 146 parts by weight of $\text{Mg}(\text{HCO}_3)_2$ hardness

3 136 parts by weight of CaSO_4 hardness

4 111 parts by weight of CaCl_2 hardness

5 164 parts by weight of $\text{Ca}(\text{NO}_3)_2$ hardness

6 120 parts by weight of MgSO_4 hardness

7 146 parts by weight of MgCl_2 hardness

8 136 parts by weight of $\text{Mg}(\text{NO}_3)_2$ hardness

The method of calculating degree of hardness will be clear from the following formula

- Hardness causing in salt in terms of CaCO_3 =

$$\frac{\text{Amount of the hardness causing salt} \times 100}{\text{Molecular weight of hardness causing salt}}$$

$$\text{Equivalents of } \text{CaCO}_3 = \frac{\left(\frac{\text{Mass of hardness}}{\text{producing substance}} \right) \times \left(\frac{\text{Chemical equivalent}}{\text{of } \text{CaCO}_3} \right)}{\left(\frac{\text{Chemical equivalent of}}{\text{hardness producing substance}} \right)}$$

UNITS OF HARDNESS.

These are 4 different units in which the hardness of water is expressed as given below

- (1) Parts per million (PPM): PPM is the number of parts of CaCO_3 equivalent hardness per 10^6 parts of water.

i.e., 1 PPM = 1 part of CaCO_3 equivalent hardness in 10^6 parts of water.

(2) Milli grams Per Litre (mg/litre): mg/L is the number of milligrams of

CaCO_3 equivalent hardness present per litre of water.

i.e., 1 mg/L = 1 mg of CaCO_3 equivalent hardness of 1 L of water.

But 1 L water weights = 1 kg of water

$$\begin{aligned}1 \text{ kg} &= 1000 \text{ gms} \\&= 1000 \times 1000 \text{ mg} \\&= 10^6 \text{ mg}\end{aligned}$$

$\therefore 1 \text{ mg/L} = 1 \text{ mg of } \text{CaCO}_3 \text{ equivalent per } 10^6 \text{ mg of water}$
 $= 1 \text{ part of } \text{CaCO}_3 \text{ equivalent per } 10^6 \text{ parts of water}$

$\therefore 1 \text{ mg/L} = 1 \text{ ppm}$

Degree Of Clark ($^{\circ}\text{Cl}$)

:

It is defined as the number of parts of CaCO_3 equivalent hardness per 70,000 parts of water.

$\therefore 1^{\circ}\text{Cl} = 1 \text{ grain of } \text{CaCO}_3 \text{ eq. hardness per gallon of}$

water. (or)

$1^{\circ}\text{Cl} = 1 \text{ part of } \text{CaCO}_3 \text{ eq. hardness per 70,000 parts of water}$

$$\therefore 1 \text{ ppm} = 0.07^{\circ}\text{Cl}$$

Degree of French (${}^{\circ}\text{Fr}$) :

${}^{\circ}\text{Fr}$ is the number of parts of CaCO_3 equivalent hardness per 10^5 parts of water.

$1 {}^{\circ}\text{Fr} = 1 \text{ part of } \text{CaCO}_3 \text{ equivalent hardness per } 10^5 \text{ parts of water}$

$$\therefore 0.1 {}^{\circ}\text{Fr} = 1 \text{ ppm}$$

Note: The hardness of water can be converted into all the four units by making use of the following interconversion formula

$$1 \text{ ppm} = 1 \text{ mg/L} = 0.07 {}^{\circ}\text{Cl} = 0.1 {}^{\circ}\text{Fr}$$

$$1 {}^{\circ}\text{Cl} = 1.43 {}^{\circ}\text{Fr} = 14.3 \text{ ppm} = 14.3 \text{ mg/L}$$

PROBLEM:

- (1) A sample of water gives an analysis 13.6 mg/L of CaSO_4 . 7.3 mg/L of $\text{Mg}(\text{HCO}_3)_2$. Calculate the total hardness and permanent hardness.

Sol:

Salt	Quantity Present (mg/L)	M.Wt	Eq. of CaCO_3
CaSO_4	13.6	136	$\frac{13.6 \times 100}{136} = 10$
$\text{Mg}(\text{HCO}_3)_2$	7.3	146	$\frac{7.3 \times 100}{146} = 5$

The Total hardness of H_2O = Temporary hardness + Permanent Hardness

$$= 5 + 10 = 15 \text{ mg/L}$$

Permanent hardness = 10 ppm (or) 10 mg/L

PROBLEM

(2) Calculate the total hardness of 1000 litre of a sample of water containing the following impurities 16.2 mg/L of $\text{Ca}(\text{HCO}_3)_2$, 11.1 mg/L of CaCl_2 , 60 mg/L of MgSO_4 and $\text{Ca}(\text{HCO}_3)_2$, 11.1 mg/L of CaCl_2 , 60 mg/L of MgSO_4 and 19 mg/L of MgCl_2 .

Sol:

Salt	Quantity Present (mg/L)	M.Wt	Eq. of CaCO_3
$\text{Ca}(\text{HCO}_3)_2$	16.2	162	$\frac{16.2 \times 100}{162} = 10$
CaCl_2	11.1	111	$\frac{11.1 \times 100}{111} = 10$
MgSO_4	60	120	$\frac{60 \times 100}{120} = 50$
MgCl_2	19	95	$\frac{19 \times 100}{95} = 20$

Total hardness of H_2O = Temporary hardness + Permanent Hardness

$$= 10 + 10 + 50 + 20 = 90 \text{ mg/L}$$

Total hardness for 1000 litres

$$= 90 \times 1000 = 90,000 \text{ mg/L}$$

PROBLEM

(3) A Sample of hard water contains the following dissolved salts per litre. $\text{CaCl}_2 = 111 \text{ mgs}$, $\text{CaSO}_4 = 1.36 \text{ mgs}$, $\text{Ca}(\text{HCO}_3)_2 = 16.2 \text{ mgs}$, $\text{Mg}(\text{HCO}_3)_2 = 14.6 \text{ mgs}$, Silica = 40 gms, Turbidity = 10 mgs.

Calculate the temporary, permanent and total hardness of water in ppm, $^{\circ}\text{cl}$ & $^{\circ}\text{Fr}$

Sol:	Salt	Quantity Present (mg/L)	M.Wt	Eq. of CaCO_3
	CaCl_2	111 mg/L	111	$\frac{111 \times 100}{111} = 100$
	CaSO_4	1.36 mg/L	136	$\frac{1.36 \times 100}{136} = 1$
	$\text{Ca}(\text{HCO}_3)_2$	16.2 mg/L	162	$\frac{16.2 \times 100}{162} = 10$
	$\text{Mg}(\text{HCO}_3)_2$	14.6 mg/L	146	$\frac{14.6 \times 100}{146} = 10$

Note: Si & Turbidity must not be considered because they do not cause hardness to water.

**Total hardness of H₂O = Hardness of Ca(HCO₃)₂ + Mg(HCO₃)₂ interms
of CaCO₃ equivalents**

$$= 10 + 10 = 20 \text{ mg/L}$$

**Permanent hardness = Hardness of CaCl₂ + CaSO₄
interms of CaCO₃ equivalents**

$$= 100 + 1 = 101 \text{ mg/L}$$

Conversion of hardness:

$$1 \text{ ppm} = 1 \text{ mg/L} = 0.07^\circ \text{cl} = 0.1^\circ \text{Fr}$$

$$\begin{aligned}\text{Total hardness of the sample of water} &= 121 \text{ ppm} = 121 \text{ mg/L} \\ &= 121 \times 0.07 = 8.47^\circ \text{cl and} \\ &= 121 \times 0.1 = 12.1^\circ \text{F}\end{aligned}$$

$$\text{Permanent hardness} = 101 \text{ mg/L, } 101 \text{ ppm, } 7.07^\circ \text{cl, } 10.1^\circ \text{ Fr}$$

$$\text{Total hardness} = 20 \text{ mg/L, } 20 \text{ ppm, } 1.4^\circ \text{cl and } 2^\circ \text{ Fr}$$

PROBLEM

(04) 1 litre of water from an underground reservoir in Tirupati town in A.P showed the following analysis for contents:

$Mg(HCO_3)_2 = 42 \text{ mg}$; $Ca(HCO_3)_2 = 146 \text{ mg}$; $CaCl_2 = 71 \text{ mg}$;

$NaOH = 40 \text{ mg}$; $MgSO_4 = 48 \text{ mg}$; Organic impurities = 100 mg;

Calculate temporary and permanent and total hardness of the sample of 10,000 lit. of water

Note: $NaOH$ & Organic impurities do not cause any hardness.

The Bicarbonate salt causes temporary hardness,

While others are responsible for Permanent hardness.

$Mg(HCO_3)_2 = 42 \text{ mg in 10,000 (or) } 42/10,000 \text{ ppm}$
Its $CaCO_3$ eq's are = $[42/10000] \times [100/146]$

Similarly for $Ca(HCO_3)_2$ it is $[146/10000] \times [100/162]$

For $CaCl_2$ it is $[71/10000] \times [100/111]$

For $MgSO_4$ it is $[48/10000] \times [100/120]$

**Temporary Hardness = [42/10000] x [100/146] + [146/10000] x
[100x162]**

Permanent Hardness = [71/10000] x [100/111] + [48/10000] x [100x120]

Total Hardness = Temporary Hardness + Permanent Hardness

$$= \dots \dots \dots + \dots \dots \dots$$

$$= \dots \dots \dots \text{ ppm}$$

PROBLEM

(5) Calculate the temporary & permanent hardness of water in °cl, containing the following dissolved salts. $\text{CaCO}_3 = 50 \text{ mg/L}$, $\text{MgCl}_2 = 9.5 \text{ Mg/L}$, $\text{CaCl}_2 = 2.2 \text{ mg/L}$ and $\text{MgSO}_4 = 12 \text{ mg/L}$

Note: CaCO_3 is an insoluble salt. It does not cause hardness. If CaCO_3 is given as H.C.S, It must be considered as $\text{Ca}(\text{HCO}_3)_2$ whose hardness is expressed in the term of CaCO_3 equivalent

Sol:

Salt	Quantity Present (mg/L)	M.Wt	Eq. of CaCO_3
CaCO_3	50 mg/L	100	50 mg/L
MgCl_2	9.5 mg/L	95	$\frac{9.5 \times 100}{95} = 10$
MgSO_4	12 mg/L	120	$\frac{12 \times 100}{12} = 10$
CaCl_2	22.2 mg/L	111	$\frac{22.2 \times 100}{111} = 20$

$$\begin{aligned}\text{Temporary Hardness Of Water} &= [\text{Hardness Of CaCO}_3] \\ &= 50 \text{ ppm}/50 \text{ mg/L} \\ &= 50 \times 0.7 = 3.5^{\circ}\text{cl}\end{aligned}$$

$$\begin{aligned}\text{Permanent Hardness Of Water} &= \text{Hardness of MgCl}_2 + \text{MgSO}_4 + \text{CaCl}_2 \\ &= 10 + 10 + 20 \\ &= 40 \text{ mg/L} \\ &= 40 \times 0.7 \\ &= 2.8^{\circ}\text{ cl}\end{aligned}$$

Problem: 100 mL of water sample has a hardness equivalent of 12.5 mL of 0.08 N MgSO₄. What is its hardness in ppm. (Mol.Wt. of MgSO₄=120)

Hardness = Hardness
12.5 mL of 0.08 N $MgSO_4$ = 100 mL
Water

$$0.08 = \frac{g}{60} = 4.8 g \text{ in } 1000 \text{ mL}$$

$$\begin{array}{c} 4.8 g \text{ in } 1000 \text{ mL} \\ \xrightarrow{\cancel{4.8 \text{ mL}}} 80 \text{ in } 12.5 \longrightarrow \frac{4.8}{1000} \times 12.5 \\ (60 \text{ mg}) \qquad \qquad \qquad = \underline{0.06 \text{ g}}. \end{array}$$

0.06 g. in 100 mL

$$80 \text{ in } 1000 \longrightarrow \frac{60}{100} \times 1000 = \underline{600 \text{ mg.}}$$

$$\begin{array}{l} \text{Hardness in} \\ \text{CaCO}_3 \text{ equivalent} \end{array} = 600 \times \frac{100}{120} = 500 \text{ ppm or mg/L}$$

X

Estimation of water Hardness

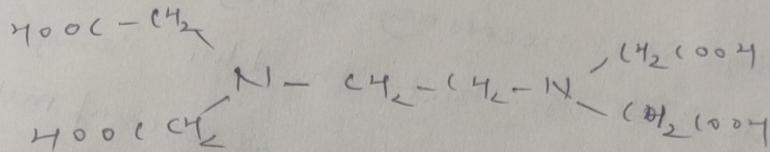
EDTA method
Soap titration method.

① EDTA method :-

Analysis is done by complexometric titration using standard EDTA [ethylene diamine tetra acetic acid] as titrant and EBT (Eriochrome Black T) as indicator

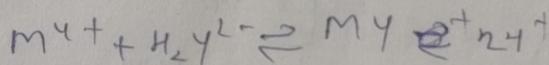
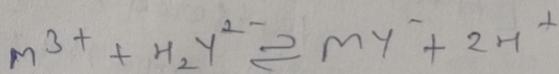
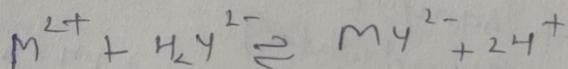
EDTA :- Tetraprotic acid $\Rightarrow H_4Y$
Limited solubility as Na_2H_2Y
salt as is used
[372.24]

as can be used as primary standard



with Ca^{2+} and Mg^{2+}

binds in 1:1

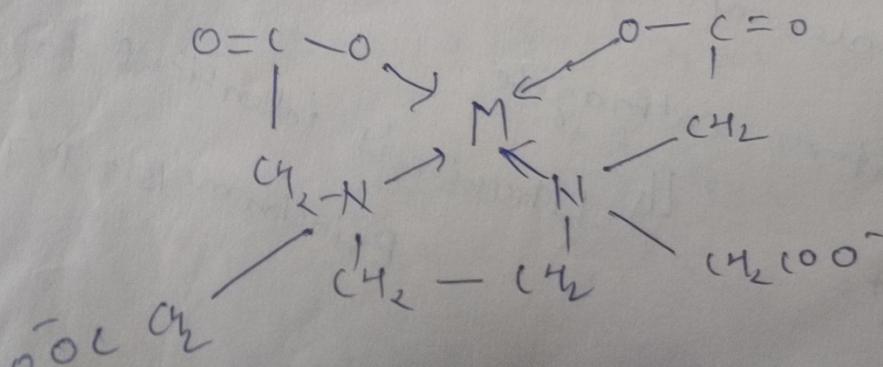


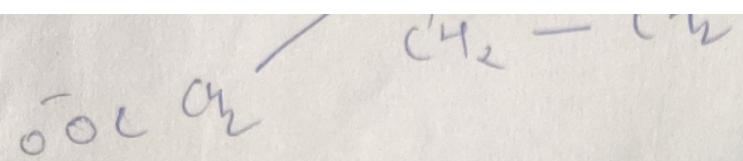
This equilibrium is markedly affected by

pH of the solution

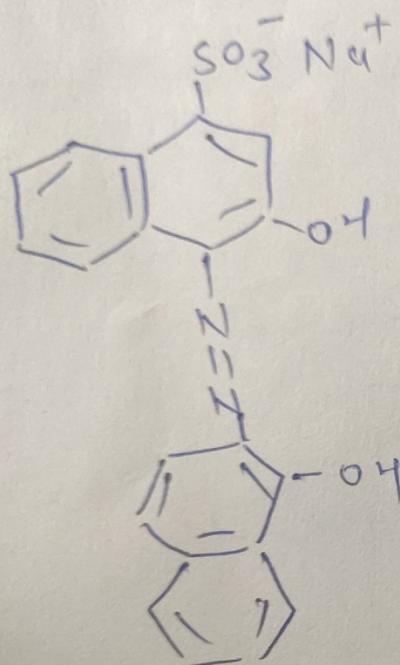
complexes are stable in basic or

slightly acidic solutions only.

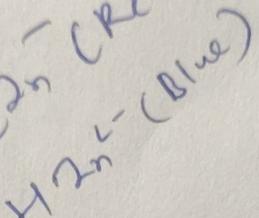
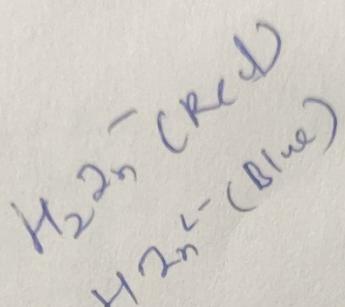




Eriochrome Black T (EBT)

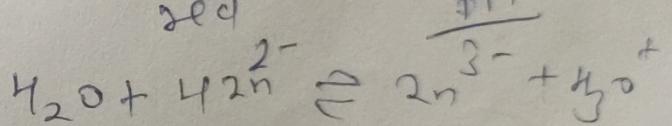
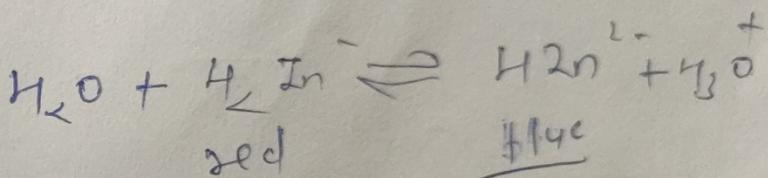


weak acid



↳ chemical name

sodium 1-(1-hydroxy-2-naphthylazo)-6-nitro-2-naphthol-4-sulphonate

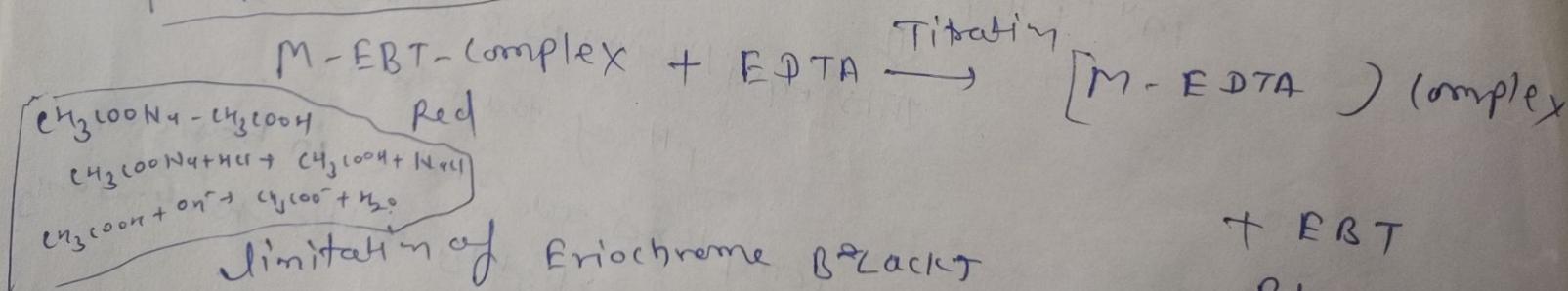
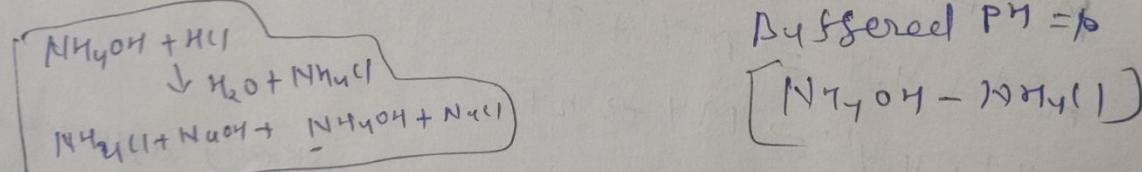
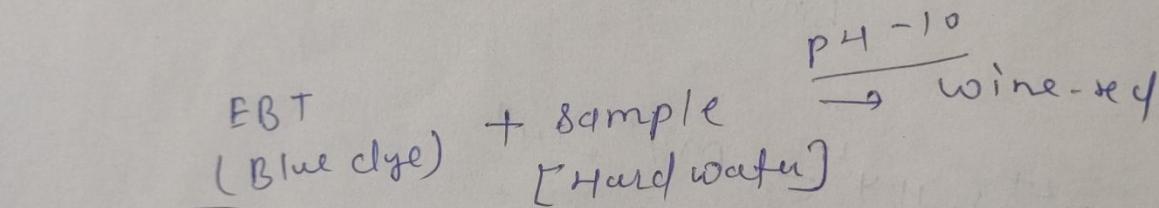


blue orange

P

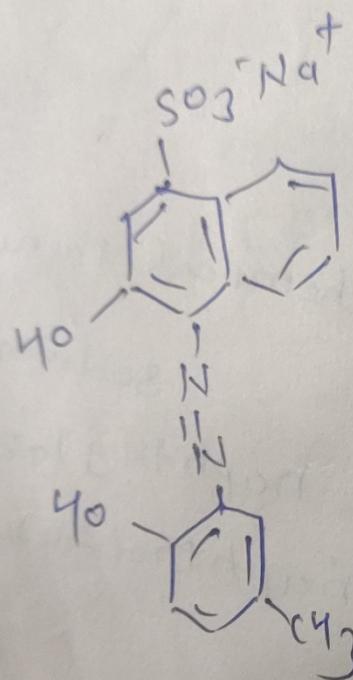
behaves as acid/base indicator as well as metal-ion indicator.

for metal ion detection it is necessary to adjust the pH on the basic side so that the blue form of species, HIn^{2-} predominates in the absence of metal ion.



that its solutions are not stable
slowly decompose on standing

Another indicator slowly dec
calmagite
ly behavior is identical to
Eriochrome Black T



Procedure:-

① Standardizing of EDTA soln:-

EDTA
↓
in burette

conical
50 mL of standard water
Prepared such a way that
it contains 1 mg.
1 mL = 1 mg of
 CaCO_3
into a 250 mL conical
flask

Standard water = 1 mg. of CaCO_3
or in 1 mL
 0.001 g. or 80
part 1 g - 100 mL conc.

molarity of standard water,

$$\frac{1}{100} = 0.01 \text{ M}$$

Add 10 mL of
buffer solution
+

Step. (i) Standardization

Std. water $\xrightarrow{\text{EDTA}}$ $M_{\text{std.}} V_1 = M_{\text{EDTA}} V_1$

$50 \text{ mL} \times 0.01 \text{ M} = M_{\text{EDTA}} V_1$

$M_{\text{EDTA}} = \frac{0.5}{V_1} M_1 = M_{\text{EDTA}}$

At the end
wine-red color changes
to blue

② Determination of total Hardness [let the volume of EDTA $\rightarrow V_2 \text{ mL}$]

50 mL of unknown water sample against EDTA

$V_{\text{unknown H}_2\text{O}} M_{\text{Unknown H}_2\text{O}} = M_{\text{EDTA}} V_2$

$V_{\text{unknown H}_2\text{O}} \times M_{\text{Unknown H}_2\text{O}} = \frac{M_2 V_2}{\text{EDTA}}$

$50 \times M_{\text{Unknown H}_2\text{O}} = \frac{M_2 V_2}{\text{EDTA}}$

$M_{\text{Unknown H}_2\text{O}} = V_2 \times \frac{0.5}{V_1} \times \frac{1}{50} = \frac{0.01 \times \frac{V_2}{V_1} M}{50}$

$$M = \frac{0.01 \frac{V_L}{V_1} \cdot g.}{100}$$

$$M = \frac{g}{1000 M \cdot ml}$$

$\checkmark M = \frac{g}{100}$

$$0.01 \frac{V_L}{V_1} \times 100 \rightarrow \text{in } 1000 \text{ mL}$$

$$0.01 \frac{V_L}{V_1} \times 100 \times 1000 \text{ mg/L}$$

$$\frac{100}{1000} \frac{V_L}{V_1} 1000 \text{ mg/L}$$

or

$$= \frac{V_L}{V_1} 1000 \text{ PPM}$$

(iii) Determination of Permanent Hardness:

~~Manganous - H_2O~~ $V_{\text{Unknown - } H_2O}$ = EDTA Take 250 mL of water $\xrightarrow{\text{Boil}}$ Reduced
~~Manganous - H_2O~~ $\times V_3$ in 50 mL to some
~~Manganous - H_2O~~ of beaker

Permanent Hardness of water.

$$= 1000 \frac{V_3}{V_1} \text{ PPM}$$

\downarrow Sintering
 \downarrow Make-up volume
 250 mL
 with distilled
 water.

(iv) Determination of temporary Hardness

Total Hardness - Permanent Hardness.

$$= 1000 \times \frac{V_2}{V_1} - 1000 \frac{V_3}{V_1}$$

\downarrow
 Titrate some of
 this water,
 against EDTA

$$\text{or } 1000 \frac{(V_L - V_3)}{V_1} \text{ PPM}$$

\downarrow
 Let the volume
 used V_3 mL

* Advantages of EDTA method :-

Determination of Hardness
 with greater accuracy, convenience and
 more rapid procedure.

Example :-

A standard hard water contains 15 g of CaCO_3 per litre, 20 ml of this required 25 ml of EDTA solutn, 100 ml of sample required 18 ml of EDTA solutn. The sample after boiling required 12 ml EDTA solutn. Calculate temporary hardness of the given sample of water in terms of ppm.

narr

Step(i) Standardization of EDTA

$$\frac{155}{1000} \quad \frac{180}{1000} \quad \frac{155}{1000} = \frac{0.83 \text{ M}}{0.15 \text{ M}}$$

$$M_{\text{std}-\text{H}_2\text{O}} V_{\text{std}-\text{H}_2\text{O}} = M_{\text{EDTA}} V_1$$

$$0.83 \times 20 = 25 \times M_{\text{EDTA}}$$

$$M_{\text{EDTA}} = \frac{0.664 \text{ M}}{0.15 \text{ M}} = 0.12 \text{ M}$$

(ii) Total hardness :-

$$M_{\text{H}_2\text{O}} V_{\text{H}_2\text{O}} = M_{\text{EDTA}} V_2$$

$$\therefore M_{\text{H}_2\text{O}} = \frac{0.12 \times 18}{100} = 0.0216 \text{ M}$$

$$0.0216 \text{ M} = 0.1195 \text{ M}$$

$$0.0216 = \frac{y}{100} \quad y = 2.16$$

$$0.1195 = \frac{g}{100} \quad g = 11.95 \text{ mg}$$

in 100 ml

11.952 mg in 100 ml

$$216^\circ \text{ ppm}$$

(iii)

Permanent Hardness

Temporary

$$216^\circ - 144^\circ = 72^\circ \text{ ppm}$$

$$M_{\text{H}_2\text{O}} V_{\text{H}_2\text{O}} = M_{\text{EDTA}} \times V_3$$

$$M_{\text{H}_2\text{O}} \times 100 = 0.664 \times 0.12 \times 12$$

$$M_{\text{H}_2\text{O}} = \frac{0.12 \times 12}{100} = 0.0144$$

$$144^\circ = \frac{y}{0.0144} \quad 0.0144 = \frac{y}{100}$$