HYDROGEN

1. INTRODUCTION:

Hydrogen is the lightest element and also the lightest gas in the periodic table.

It is the lightest non-metal. It is the simplest element in periodic table having only 16 1p & no neutron.

Electronic configuration of H is 1s1

Number of $e^- = 1$ Number of orbital = 1 Number of shells = 1 Number of subshell =1

It is discovered by Henry Cavendish & it was called inflammable element.

The name hydrogen was given by Lavoisier





Hydro (Water)

Gene (Producer) or generator

means according to Lavoisier the gas which produce water on burning with oxygen.

$$H_2 + \frac{1}{2}O_2 \to H_2O$$

Order of Abundance of H:

Hydrogen is the most abundant element in the universe (70%) of the total mass of the universe.)

Order of abundance of H:

Universe > Sun atm > Earth

The planet Jupiter & Saturn consist mainly of H₂. Similarly about half the mass of the sun & some other stars is made up of hydrogen.

In Sun's atomsphere & in universe, It is found in atomic form. While in earth it is generally found in molecular form. At Sun, the stratosphere is made up of H (atomic hydrogen) & they undergoes fusion & converted into He nuclei & this reaction is exothermic so lot of amount of energy is liberated.

It is the ninth element on earth in order of abundance.

Earth does not posses enough gravitational force to retain live hydrogen molecule i.e. why it is not found in earth atomosphere in atomic form.

Hydrogen is the most reactive elements in atomic form but it is less reactive in molecular form because of very high bond dissociation energy due to 1s -1s overlapping.

2. POSITION OF H, IN PERIODIC TABLE :

Hydrogen is the tirst element of Periodic table but still it could not be assigned a proper position either in Mendeleef periodic table or in Modern periodic table because of following reasons.

It may kept in 19/IA or 17th/VIIA group due to following reason.

Resembeles with 1st/IA Alkali metals.

(i) Electronic configuration: Like alkali metals hydrogen also has only one electron in outer most shell.

$$H = 1s^{1}$$
 $K = 4s^{1}$
 $Li = 2s^{1}$ $Rb = 5s^{1}$ ns $Na = 3s^{1}$

i) Electropositive characters: Like alkali metals hydrogen also have the tendency to loose one electron to form cation.

e.g.
$$Na \rightarrow Na^+ + e^ H \rightarrow H^+ + e^-$$

(Proton)

(iii) Oxidation numbers: Like alkali metals hydrogen can also exhibit the oxidation number of +1 in most of its compound.

e.g. $Na^+Cl^ K^+Cl$ H^+Cl^- = +1

(iv) Reaction with electronegative elements (non-metals): Like alkali metals H also reacts with exygen, Sulphur, Halogens to form oxides, sulphides and halides respectively.

$$O \xrightarrow{gives} H_2O, \text{ like Na}_2O, K_2O$$

$$S \xrightarrow{gives} H_2S$$
, like Na_2S , K_2S

with halogen \xrightarrow{gives} HX, like NaX, KX

(v) Liberation at Cathode:

$$2Na^+ Cl$$
 electrolysis $2Na$ + Cl_2 (liberated at cathode) (liberated at anode)

 (H_{a})

$$2H^+$$
 Cl^- electrolysis

Cl₂)

like alkali metals H also get liberated at cathode on electrolysis.

(vi) Reducing nature: Like alkali metals H₂ also have reducing nature.

$$CuO + H_2 \longrightarrow Cu + H_2O$$

Due to resembling of these properties with alkali metals H can be placed in IA/group /alkali metals group.

Resembles with halogen/ VIIA/17th

(i) Electronic configuration:

Electronic configuration of halogen
$$H \longrightarrow 1s^1$$

both these requires one electron to attain the stable configuration of their nearest inert gas.

$$H \longrightarrow 1s^1$$
 Recess than He $(1s^2)$ F $\longrightarrow 2p^5$ Recess than Ne $(2p^6)$ C $\longrightarrow 3p^5$ Recess than Ar $(3p^6)$

(ii) Electronegative character: Both halogen & Hydrogen has the tendency to gain one e-

$$F + e^{-} \longrightarrow F^{-}$$

$$H + e^{-} \longrightarrow H^{-}$$

i.e. why both will act as electronegative species.

- (iii) lonization potential: Ionization potential of hydrogen is almost similar to halogen.
- (iv) Oxidation number: Both halogen & hydrogen can exahibit –1 oxidation number.

Hydrogen in metal hydride shows -1 oxidation state.

$$NaH \longrightarrow Na^{+} + H^{-}$$

$$NaCl \longrightarrow Na^{+} + Cl^{-}$$

- (v) Diatomic molecule: Both halogen & hydrogen have the tendency to exist as diatomic molecule. eg. F_2 , Cl_2 Br_2 , I_2 & H_2 .
- (vi) Liberation at anode: Whenever Alkali metals halides & hydrides undergoes electrolysis, both halogen and hydrogen will liberate at anode.

$$2Na^+ Cl^- \longrightarrow 2Na + Cl_2$$
 (anode)
 $2Na^+ H^- \longrightarrow 2Na + H_2$ (anode)

(vii) Reaction with highly electropositive metals: Hydrogen reacts with highly electropositive element (i.e. s-block elements) & forms their hydrides like their halides.

$$\begin{array}{cccc} 2 \text{Na} & + & \text{H}_{_2} & \longrightarrow & 2 \text{NaH} \\ 2 \text{Na} & + & \text{Cl}_{_2} & \longrightarrow & 2 \text{NaCl} \end{array}$$

This reaction shows oxidising character of hydrogen.

(viii) Formation of covalent compound: Both Hydrogen & halogen on reaction with non metals to form covalent compounds like.

Conclusion : The position of hydrogen is still in controversy & no proper position is assigned to H in periodic table i.e. why it is also called notorious or rogue elements.

3. ISOTOPES OF HYDROGEN:

There are 3 isotopes of H_2

	Protium or ordinary hydrogen	Deuterium/ Heavy hydrogen	Tritium	
Symbol	₁ H ¹ H ₂	1H2/P2	₁ H ³ / ₁ T ³ T ₂	
Property	Protium or ordinary hydrogen	Deuterium or heavy hydrogen	Tritium	
Atomic number	1	1	1	
Mass number	1	2	3	
Exact atomic mass	1,008123	2.0142	3.0170	
Symbol	H	$_{1}^{2}\mathrm{H}$ or $_{1}^{2}\mathrm{D}$	${}^3_1\mathrm{H}$ or ${}^3_1\mathrm{T}$	
Molecular formula	H	D_2	$T_{_2}$	
No. of protons in			2	
the nucleus		1	1	
No. of neutrons in				
the nucelus	Nil	1	2	
No. of electron	1	1	1	
Electronic configu.	1s ¹	$1s^1$	$1s^1$	
Relative abundance	99.984%	0.016%	10^{-15} %	
Stability	Stable	Stable	Unstable(Radioactive)	

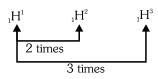
Properties	H ₂	D ₂	T ₂
MA	- 259°C	- 254.3°C	- 252.4°C
B.P.	- 252.6°C	- 249.3°C	- 248.0°C
Bond length (H- H)	74 pm	74 pm	74 pm
Bond energy (H—H)	436.0 KJ mol ⁻¹	443.3 KJ mol ⁻¹	446.9 KJ mol ⁻¹
Heat of fusion & vaporisation	Minimum	H < D < T	Maximum

ISOTOPIC EFFECT: 4.

The effect which can change the physical & chemical properties of isotopes is called isotopic effect. It is because of difference in mass.

In isotopic effect maximum changes occurs in physical properties like melting point, boiling point, bond energy while minimum changes occurs in chemical properties like state of chemical reaction etc.

Imp. Isotopic effect is found only in hydrogen isotopes. Because there is large difference in mass.



While in other isotopes like $_{7}^{}N^{14}$ $_{7}^{}N^{15}$ only a fractional mass is increased.

 ${}_{6}C^{14}$ ${}_{6}C^{13}$ ${}_{6}C^{12}$ \longrightarrow Here also only a fraction is increased

But in case of hydrogen mass increased to 2 to 3 times in their isotopes.

- Isotopic effect is found in: Q.
 - (1) H
- (2) N
- (3) C
- (4) All

Η

1 0

Atomic hydrogen

Ans. (1)

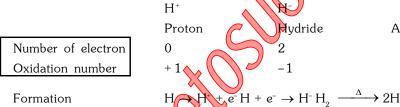
- Q. Which of the following reaction is fast & why?
 - (i) $CH_4 + Cl_2 \longrightarrow CH_3Cl + HCl$
 - (ii) $CD_4 + Cl_2 \longrightarrow CD_3Cl + DCl$

Ans. (i) because C-H bond energy is less in comparison to C-D bond energy.

Different forms of Hydrogen:

Based on oxidation Number. (a)

There are three types of hydrogen



Formation

Note: In the aqueous state proton (H) exist as H+ (H₂O),

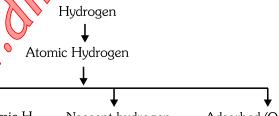
Where n is a large number.

If
$$n = 1$$
 $H_3^{0^+}$ $n = 2$ $H_1^+(H_2^-O)_2$

- Q. In the equeous solution hydrogen ion exist as
 - (1) H₂O⁺
- (2) H (H₂O)₂
- (3) $H^{+}(H_{0}O)_{-}$
- (4) All

(4) Ans.

> (b) Based on reactivity:



Simple atomic H

Nascent hydrogen

Adsorbed/Occofluded

Atomic hydrogen :

Simple atomic hydrogen - It is formed by simple dissociation of hydrogen. (i)

Favourable comdition - Favourable condition are high temp & low pressure.

(ii) Nascent hydrogen – Hydrogen at the moment of its birth it called nascent hydrogen means which torms at the instant is known as Nascent hydrogen.

It is formed only by some specific chemical reaction.

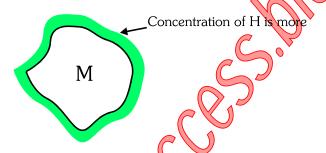
(a) Acid + Metals

$$Zn + H_2SO_4 \longrightarrow ZnSO_4 + 2H$$

- (b) Base + element $2NaOH + Be \longrightarrow Na_9BeO_9 + 2H$
- (c) $C_{2}H_{5}OH + Alkali metal$

$$C_2H_5OH + Na \longrightarrow C_2H_5ONa + H$$

(iii) Adsorbed/Occluded hydrogens



Adsorbed H is hydrogen present at the outer surface of metal.

Occlusion - The property of metal to adsorb any gas is called occlusion.

Reactivity order

Atomic hydrogen > Nascent hydrogen > Molecular hydrogen

(iii) Based on Nuclear spin (Nuclear isomers)



- (a) Ortho hydrogen— The molecular form of hydrogen having same spin of proton is called ortho hydrogen.
- (b) Para hydrogen The molecular form of hydrogen having opposite spin of proton is called para hydrgen. In ortho hydrogen spin of proton is same, so they will repel each other & because of this repulsion, internal energy of ortho hydrogen increases. So ortho hydrogen has more internal energy.

Stability of ortho & para hydrogen

Stability of ortho & para hydrogen depends upon temperature condition.

At low temp: para hydrogen is more stable than ortho hydrogen while at high temp ortho hydrogen is more stable than para hydrogen.

$$P_{H_2} \xrightarrow{T\uparrow} O_{H_2}$$

	Ortho	Para
At 25 C	75%	25%
At -253 C/20K	0	100%

Imp. Note: (i) We can obtain 100% pure para hydrogen at low temp but can't ortho because at high temp parahydrogen will dissociate into atomic hydrogen.

(ii) Ortho & Para hydrogen differs only in physical properties but have same chemical properites.

Method of preparation:

(A) From acids: The metal which are placed about H_2 in electrochemical series react with dil acids they liberate H_2 .

e.g. Fe +
$$H_2SO_4 \rightarrow FeSO_4 + H_2$$
 (dil)
 $Cu + H_2SO_4 \rightarrow$ (No reaction)

Lab preparation: When impure Zn reacts with dil H2SO4 it forms H2

$$Zn$$
 + $H_2SO_4 \rightarrow ZnSO_4 + H_2$ (impure) (dil)

Q. Why we use impure Zn.

Ans. Because the rate of reaction with pure Zn is very slow.

2. By alkalies: Only (Be, Zn, Al, Sn, Pb, Si) (Amphoteric metal) react with boiling NaOH or KOH they evolve H₂.

$$Zn + 2NaOH \rightarrow Na_2ZnO_2 + H_2\uparrow$$
 (sodium zincate)

2Al + 2NaOH +
$$2H_2O \longrightarrow NaAlO_2 + 3H_2\uparrow$$

(sodium meta aluminate)

$$Sn + 2NaOH + H_2O \longrightarrow Na_2SnO_3 + 2H_2 \uparrow \uparrow$$

(sodium stannate)

Pb + 2NaOH +
$$H_2O \longrightarrow Na_2PbO_3 + 2H_2$$

(sodium plumbate)

$$Si + 2NaOH + H_2O \longrightarrow Na_2SiO_3 + 2H_2\uparrow$$

(sodium silicate)

Be + 2NaOH
$$\longrightarrow$$
 Na₂BeO₂ + $\stackrel{\text{(s)}}{\text{(s)}}$

(sodium beryliate)

3. From water:

All the metals which are placed above than H₂ when react with water the evolve H₂.

$$Zn + H_2O \longrightarrow ZnO + H_2\uparrow$$

Three type of water is used

(i) Cold water: The temperature of cold water is 7 to 25 C this water is used for highly reactive metals.

Such as Lack, Bay Sr, Ca, Na, means alkali metals of alkalic earth metals.

The reaction with alkali metals are vigorous to minimum the rate of reaction these metals are used in the form of amalgam.

$$2Na(Hg) + H_2O \longrightarrow 2NaOH + H_2\uparrow$$

(ii) Hot water: The temperature of hot water is 25 C to 90 C. This water is used for reactive metals, such as Mg, Al, Mn, Zn, Cr.

(iii) Steam: The temperature of steam is more than 100 C. This form of water is used for very less reactive metals like Fe, Cd, Co, Ni, Sn, Pb.

Condition for best yield of H₂.

- (i) Cold water \longrightarrow With highly reactive metals.
- (ii) Hot water → With reactive metals
- (iii) Steam \longrightarrow With less reactive metals.

4. On Ionic hydride:

Whenever ionic hydride reacts with water then form H_2 .

$$CaH_2 + 2H_2O \longrightarrow Ca(OH)_2 + H_2$$

(Hydrolith)
 $NaBH_4 + H_2O \longrightarrow NaBO_2 + 4H_2$

Method to prepare pure hydrogen :

Electrolysis of water: To prepare pure hydrogen we use impure water (i.e. having 15-20% solution of or acid)

$$4H_{\circ}O \rightleftharpoons 4H^{+} + 4OH^{-}$$

at cathode
$$4H^+ + 4e^- \longrightarrow 2H_2$$

at anode
$$4OH^- \longrightarrow 2H_2O + O_2 + 4e^-$$

The SO₄-2 or K⁺ ion present in acid or alkali does not move towards anode or eathore as their discharge potential is higher than of OH- ions or H+ ions rescrectively.

2. By reaction of Magnesium with dil. H,SO4:

$$Mg + H_2SO_4 \longrightarrow MgSO_4 + H_2$$

3. B reaction of NaH with water :

$$NaH + H_{2}O \longrightarrow NaOH + H_{2}$$

4. Uyeno method: This method is used for military purpose

$$2Al + 2KOH + 2H_2O \longrightarrow 2KAlO_2 + 3H_2\uparrow$$

(Potassium meta aluminate)

Because by this reaction we can prepare H₂ in a rapid namer.

Commercial or industrial method to prepare H_2 :

Lane process: Steam is passes over hot in the converts into Fe₃O₄ & H₉. (i)

$$3Fe + 4H_2O \longrightarrow Fe_3O_4 + 4H_2$$

Iron is regenrated by reducing $\mathrm{Fe_3O_4}$ into Fe by water gas (CO + $\mathrm{H_2}$)

$$2Fe_3O_4 + 4CO + 4H_2O$$

 $Fe_3O_4 + 2CO + 2H_2O$
 $3Fe + 2CO_2 + 2H_2O$

$$Fe_3O_4 + 2CO + 2H_2 \longrightarrow 3Fe + 2CO_2 + 2H_2O$$

This process is a continuous process.

Bosch Process: In this process initially steam is passed over red hot coke then water gas is formed. (ii)

$$C + H_2O \longrightarrow CO + H_2$$

Then water gas is mixed with more stem in presence of Fe₂O₃/Cr₂O₃ then CO will convert into O₂ & we can obtain more 🅀

$$CO + H_2 \rightarrow H_2O \longrightarrow CO_2 + 2H_2$$

(iii)

From Natural gas:
$$\begin{array}{c} CO + H_2 \cap H_2O \longrightarrow CO_2 + 2H_2 \\ \hline \text{From Natural gas}: \\ C_n H_{3n+2} + nH_2O \longrightarrow nCO + (2n+1)H_2 \end{array}$$

- PHYSICAL PROPERTIES OF HYDROGEN: 5.
 - Hydrogen is a lightest, colourless, odourless and tasteless gas. It is sparingly soluble in water. It is inflammable and less reactive gas.
 - (-259.2°C) and b.p. (-252°C) are very low indicating less intermolecular attraction. Due to low to. Tiquid hydrogen is used as a cryogenic fluid (to produce low temperature).
 - HHH bond energy [$104~{
 m Kcal~mol^{-1}}$] and $436~{
 m KJ/m}$
 - $^{\circ}$ H H bond length [74 pm] so $\mathrm{H_{2}}$ is less reactive and require high temp for reaction.

Chemical properties of H₂:

- H_o is neutral in nature i.e. why it does not react with acids & bases. (i)
- (ii) Less reactive: Hydrogen is very less reactive in nature because of very high bond dissociation energy.

(iii) Combustible nature: H_2 is highly combustible in nature & it burns with oxygen or air with pale blue flame to give water.

$$H_2 + \frac{1}{2}O_2 \longrightarrow H_2O$$

(iv) Reaction with highly electropositive metals :

IA & IIA group elements are called highly electropsitive metal. Whenever H reacts with these metals they form ionic hydrides.

$$\begin{array}{ccc} \mathrm{2Na} \ + \ \mathrm{H_{2}} & \longrightarrow \ \mathrm{2NaH} \\ \mathrm{CO} \ + \ \mathrm{H_{2}} & \longrightarrow \ \mathrm{CaH_{2}} \end{array}$$

- (v) Reaction with non-metal:
 - (a) Reaction with halogen

order of reactivity of halogen with hydrogen

$$F_2 > Cl_2 > Br_2 > I_2$$

(b) Reaction with sulphur

$$H_2 + S \longrightarrow H_2S$$

(c) With nitrogen - (Haber process)

$$N_2 + 3H_2 \xrightarrow{\text{Fe, Mo, 200atm}} 2NH$$

(vi) Reducing nature:

$$CuO + H_2 \longrightarrow Cu + H_2O$$

(vii) Reaction with carbon mono oxide:

(viii) Hydrogenation of unsaturated hydrocarbon:

$$C_{2}H_{2} + 2H_{2} \xrightarrow{C_{2}H_{6}}$$

$$C_{2}H_{4} + H_{2} \xrightarrow{A73K} C_{2}H_{6}$$

When unsaturated hydrocarbon having C = C or $C \equiv C$ reacts with hydrogens in the presence of Ni/Pd/Pt forms saturated hydrocarbons.

Imp. Hydrogenation of vegetable oil.

Vegetable oils are also called polyunsaturated oils because they contain many C = C bond. When these oil are exposed to all for a long time then double bond will get oxidized and the oil becomes. Rancid (having fowl. smell or unpleasant test) in nature. So to avoid this vegetable oil are converted into edible fats (Vanaspati Ghee)

Vegetable oils +
$$H_2 \xrightarrow{\text{Ni}}$$
 edible fats solid (liquid)

This whole process is known as hydrogenation or hardening of oil.

- 6. USES OF HYDROGEN:
 - Hydrogenation of vegetable oil to form solid fats i.e. vanaspati ghee.

In liquid form as a rocket fuel. (Liquid H_2 + Liquid O_2)

- 3. In a air ship of balloons as a mixture of Hydrogen & Helium [15% $m H_2$ + 85% He]
- 4. Formation of different compounds.

Like \rightarrow NH₃, (Haber process) alkane, alcohol and other hydrocarbon

Hydrides: The compounds of hydrogen with different elements are called hydrides.

These are of three types:

(1) Ionic/Salt like/Saline hydrides :

Compounds of hydrogen with s-block elements except berylium & magnesium are called ionic hydride

LiH, NaH, KH, RbH, CsH, CaH
$$_2$$
, SrH $_2$, BaH $_2$

BeH₂, MgH₂ are covalent polymeric hydride.

- u Structure of these hydrides are similar to rock salt, so they are also called salt like/saline hydrides
- u Down the group size↑ Lattice energy↓ stability↓ Melting point↓ Boiling point↓
- u On electrolysis of these hydrides, hydrogen is liberated at anode.
- u On reaction with water these hydrides will form hydrogen

$$NaH + H_2O \longrightarrow NaOH + H_2$$

u These hydrides forms complex hydrides which are very good reducing agent

$$4LiH + AlCl_3 \longrightarrow LiAlH_4 + 3LiCl$$

 $NaBH_4 \longrightarrow Sodium borohydride$

 $LiAlH_4 \longrightarrow Lithium$ aluminium hydride.

(2) Mettalic / Interstitial hydrides :

They are the compounds of d & f-block elements.

In these hydrides hydrogen occupies interstitial sites present in metallic lattice, so they are called interstitial hydrides.



u Properties of these hydrides are similar to parent metals, so they are also known as metallic hydrides.

u These hydrides are non. stochiometric in nature (i.e. having variable composition)

$$ZrHx (x = 1.3 - 1.75)$$

$$TiHx (x = 1.8 - 2)$$

u Metals of group 7,8,9 donot form any hydrides so this particular part of periodic table is known as hydride gap.

(3) Covalent/Molecules hydrides

They are the compounds of hydrogen with p-block elements CH_4 , NH_3 , H_2O , HF, etc.

These hydrides exist as molecules, so they are also known as molecular hydrides. There hydrides are non-conductor of electricity.

Nomenclature - element Suffix (ane)

$$NH_3 \longrightarrow Azane$$

$$H_0O \longrightarrow Oxidane$$

These hydrides are again divides into 3 categories.

(a) Electron deficient hydrides:

u They are the hydrides of group 13 elements.

 BH_3 , AIH_3 , GaH_3 - In these hydrides central element does not have complete octet. i.e. why they are called electron deficient compounds.

(b) Electron precise hydrides - They are the hydrides of group 14 element.

In these type of hydrides central elements has 8e- in their outer most shell.

Electron rich hydrides: These are the hydrides of group 15, 16, 17

e.g.
$$\ddot{N}H_3$$
, $H_2\ddot{O}$, $H\ddot{F}$: etc.

In these hydrides lone pair are present on central dement which can be given to others. So they are called electron rich hydrides.

Water (H₂O)

Properties of pure water :

(a) Physical properties:

(i) Pure water is colourless, tasteless and odourless. It gives bluish tinge in thick layers. (ii) It freezes at 0 C and boils at 100 C. (iii) Its maximum density is 1.00 at 4 C. (iv) It is a polar molecule and has V-shaped structure. The bond angle is 104.5. (v) It has a high dielectric constant. The polar character of water makes it an excellent solvent for polar and ionic substances. (vi) It is a poor conductor of electricity. (vii) It has the tendency to associate. It exists in the liquid state not as a single H₂O molecule but as associated molecules through-hydrogen bonding. The existence of hydrogen bonding is responsible for high values of specific heat, the latent heat of fusion and latent heat of vaporisation.

(b) Chemical properties:

(i) Water is neutral in nature. pH of the pure water is 7. It is a weak electrolyte and feebly ionises into H⁺ and OH⁻ ions.

$$H_0O \rightleftharpoons H^+ + OH^-$$

In pure water $[H^+] = [OH^-] = 10^{-7}$ at 25 C.

(ii) With metals: it reacts with active metals and evolves hydrogen. The reaction is exothermic in the case of alkali and alkaline earth metals.

$$2Na + 2H_2O \longrightarrow 2NaOH + H_2$$

 $Ca + 2H_2O \longrightarrow Ca(OH)_2 + H_2$

(iii) Reaction with non metals :

Chlorine decomposes cold water forming HCl and HClO.

$$Cl_2 + H_2O \longrightarrow HCl$$
 + HCO
(Hypothbrius acid

When steam is passed over red hot coke (1000 C), water gas is formed.

$$C + H_2O \rightarrow \underbrace{CO + H_2}_{Water gas}$$

(iv) Action on nonmetalic oxides: Acidic oxides combine with water to form acids.

$$CO_{2} + H_{2}O \longrightarrow H_{2}CO_{3}$$
 (Carbonic acid)
$$SO_{2} + H_{2}O \longrightarrow H_{2}SO_{3}$$
 (Sulphurous acid)
$$SO_{3} + H_{2}O \longrightarrow H_{2}SO_{4}$$
 (Sulphuric acid)
$$P_{2}O_{5} + 3H_{2}O \longrightarrow 2H_{3}PO_{4}$$
 (Orthophosphoric acid)
$$N_{2}O_{5} + H_{2}O \longrightarrow 2HNO_{3}$$
 (Nitric acid)
$$C_{1}O_{7} + H_{2}O \longrightarrow 2HCIO_{4}$$
 (Perchloric acid)

(v) Action on metallic oxides: basic oxides combine with water to form alkalies.

$$Na_2O + H_2O \longrightarrow 2NaOH$$
 $CaO + H_2O \longrightarrow Ca(OH)_2$

Action on hydrides, carbides, nitrides, phosphides: Water decomposes these compounds with liberation of hydrogen, acetylene (or methane), ammonia, phosphine respectively.

$$CaH_2 + 2H_2O \longrightarrow Ca(OH)_2 + 2H_2$$

 $CaC_2 + 2H_2O \longrightarrow Ca(OH)_2 + C_2H_2$

$$Al_4C_3 + 12H_2O \longrightarrow 4Al(OH)_3 + 3CH_4$$
 $Mg_3N_2 + 6H_2O \longrightarrow 3Mg(OH)_2 + 2NH_3$
 $Ca_3P_2 + 6H_2O \longrightarrow 3Ca(OH)_2 + 2PH_3$

(vii) **Hydrolysis**: Many salts specially the salts of strong bases with weak acids, weak bases with strong acids and weak bases with weak acids undergo hydrolysis with water.

$$CH_3COONa + H_2O \rightleftharpoons CH_3COOH + NaOH$$

 $CH_3COONH_4 + H_2O \rightleftharpoons CH_3COOH + NH_4OH$

Halides of nonmetals are decomposed by water.

$$PCl_5 + 4H_2O \longrightarrow H_3PO_4 + 5HCl$$

 $PCl_3 + 3H_2O \longrightarrow H_3PO_3 + 3HCl$

Hard and soft water

A water is said to be a soft water if it produces sufficient lather with the soap and water is described as being hard if it forms an insoluble scum before it forms a lather with soap. The hardness of natural water is generally caused by presence of bicarbonates, chlorides and sulphates of cardium and magnesium but infact soluble salts that form a scum with soap cause hardness.

$$\begin{aligned} &\text{Ca$^{2+}$(aq) + 2C$_{17}$H$_{35}$COO$^-$ (aq) } \longrightarrow &\text{(C$_{17}$H$_{35}$COO)$_2$Ca} \\ &\text{Mg$^{2+}$ (aq) + 2C$_{17}$H$_{35}$COO$^-$ (aq) } \longrightarrow &\text{C$_{17}$H$_{35}$COO]$_2$Mg} \\ &\text{Anion of soap} \end{aligned}$$

Soap will not produce lather with water until all the calcium and magnesium ions have been precipitated. Hard water thus wastes soap. Hardness of water is of two types:

- (a) Temporary hardness
- (b) Permanent hardness
- (a) Temporary hardness: This is due to the presence of bicarbonates of calcium and magnesium.

Temporary hardness in water is easily removed by boiling, as the bicarbonates decompose readily and the insoluble carbonates are precipitated.

$$Ca(HCO_3)_2 \xrightarrow{\text{Roil}} CaCO_3 + H_2O + CO_2$$

$$Mg(HCO_3)_2 \xrightarrow{\text{Roil}} MgCO_3 + H_2O + CO_2$$

$$(Inso lub le)$$

Temporary hardness can also be removed by Clark's process which involves the addition of slaked lime [Ca(OH)₂].

$$Ca(HCO_3)_2 + Ca(OH)_2 \longrightarrow 2CaCO_{(lnso libble)} + 2H_2O_3$$

this essential to add only the calculated amount of Ca(OH), because excess will cause artificial hardness.

Permanent hardness: Permanent hardness is introduced when water passes over rocks containing the sulphates or chlorides of both of calcium and magnesium. This type of hardness cannot be removed by boiling or by the addition of slaked lime.

The various water softeners are :

$$CaCl_2 + NaCO_3 = CaCO_3 + 2NaCl$$

$$CaSO_4 + Na_2CO_3 = CaCO_3 + Na_2SO_4$$

$$Ca(HCO_3)_2 + Na_2CO_3 = CaCO_4 + 2NaHCO_3$$
(So bub le)

In place of sodium carbonate, caustic soda or sodium phosphate can also be used

$$MgCl_2 + 2NaOH \longrightarrow Mg(OH)_2 + 2NaCl_{(Inso lub le)}$$

$$3MgSO_4 + 2Na_3PO_4 \longrightarrow Mg_3(PO_4)_2 + 3Na_2SO_4$$

(ii) Calgon: The complex salt of metaphosphoric acid, sodium hexametaphosphate (NaPO₃)₆, is known as calgon. It is represented as Na₂[Na₄(PO₃)₆]. Calcium and magnesium salts present in hard water react with calgon to give complex salts.

$$2CaSO4 + Na2[Na4(PO3)6] \longrightarrow Na2[Ca2(PO3)6] + 2Na2SO4$$
$$2MgSO4 + Na2[Na4(PO3)6] \longrightarrow Na2[Mg2(PO3)6] + 2Na2SO4$$

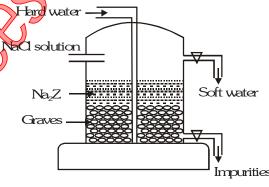
(iii) Permutit process: Permutit is hydrated sodium alumino silicate

$$[Na_2Al_2Si_2O_8. xH_2O]$$

$$[Na_2O.Al_2O_3.2SiO_2.xH_2O]$$

Permutit also known as sodium zeolite (Na₂Z), means Zeolite is Al₂Si₂O₈·xH₂O.

In this process when hard water is poured into chamber, it may contain organic impurities like plant. This impurities can be removed by gravel.



On moving upwards hard water will react with Na_2Z during this reaction Na^+ ions of Na_2Z will replace Mg^{+2} & Ca^{-2} of impurities.

$$Na_2Z + CaC_2 \longrightarrow 2NaCl + CaZ$$

NaCl is dissolved in water & water becomes soft.

This soft water is fit for washing purpose but not fit for drinking purpose.

After some time when Na_2Z is completely converted into CaZ. Process is stopped and for pregeneration of Na_2Z . We use NaCl solution.

$$2$$
NaCl + CaZ \longrightarrow CaCl₂ + Na₂Z

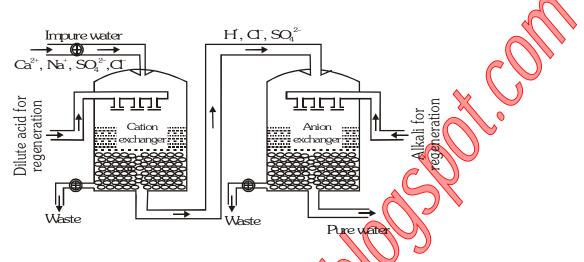
for exchange resin: By this process we can remove both cation & anion of hardness.

This process contain two chamber.

- (a) Cation exchange resin: This resin contains granular insoluble organic acid having giant molecules with -COOH group.
- (b) Anion exchange resin: This resin contain giant organic molecules with basic groups derived from amines.

Process : When hard water is poured into first chamber the cation of hardness $(Mg^{+2}. Ca^{+2})$ removed by H^+ ions of organic acid.

$$\begin{split} 2\text{R-COO}^-\text{H}^+ + \text{CaCl}_2 & \longrightarrow \text{(R-COO)}_2\text{Ca} + 2\text{H}^+ + 2\text{Cl}^- \\ + \text{MgSO}_4 & \longrightarrow \text{(R-COO)}_2\text{Mg} + 2\text{H}^+ + \text{SO}_4^{-2} \end{split}$$



This water becomes soften but not used for drinking burpose because this water contain the impurity of acid. To remove anion of hardness this acidic water then passed through another bed containing anion exchanger. This exchanger removes anion like Cl^- , SO_4^{-2} & HCO_3^{-1} .

$$R-NH_3^+OH^- + H^+ + Cl^- \longrightarrow R-NH_3^+Cl^- H^+OH^-$$

This water is free from impurities & can be used for drinking purpose.

After some times when both resin gets expansived process is stopped.

Regenration of resin:

(i) Cation exchange resin: We use dil acid.

$$2H^+Cl^- + (R-COO)_2Ca \longrightarrow 2R^-COO^-H^+ + CaCl_2$$

(ii) Anion exchange resin We use dil NaOH solution

$$R-NH_3^+Cl \longrightarrow Na^+OH^- \longrightarrow Na^+Cl^- + R-NH_3^+OH^-$$

Heavy water (D₂O)

Method of preparation:

Repeated electrolysis of H_2O : On electrolysis of water (impure) H_2O dissociate into H^+ & OH^- while a fractional part of D_2O will dissociate into D^+ & OD^-

$$H_2O \rightleftharpoons H^+ + OH^ D_2O \rightleftharpoons D^- + OD^-$$

 D^+ & OD^- due to more mass have less mobility i.e. why H^+ & OH^- will move towards cathode & anode respectively while D^+ & OD^+ will be in solution.

This process will repeated for six time.

Properties of Heavy water:

Physical properties: (a) Heavy water is a colourles, odourless and tasteless mobile liquid, (b) Nearly all the physical constants are higher than the corresponding values of ordinary water.

Chemical properties: Heavy water is chemically similar to ordinary water. However, D_2O reacts more slowly than H_2O in chemical reactions.

(a) Action of metals: D2O reacts with alkali and alkaline earth metals liberates heavy hydrogen.

$$2D_2O + 2Na \longrightarrow 2NaOD \atop Sodium deuteroxide + D_2$$

$$2D_2O + Ca \longrightarrow Ca(OD)_2 + D_2$$

Action with metallic oxides: D2O reacts slowly with basic oxides to form heavy alkalies (b)

$$Na_2O + D_2O \longrightarrow 2NaOD$$

$$CaO + D_{2}O \longrightarrow Ca(OD)_{2}$$

Action with nonmetallic oxides: D₂O reacts slowly with acidic oxides to form deutero acids. (c)

$$N_2O_5 + D_2O \longrightarrow 2DNO_3$$
Deutero nitric acid

$${\rm P_2O_5} \, + \, 3{\rm D_2O} \longrightarrow \\ \begin{array}{c} 2{\rm D_3PO_4} \\ {\rm Deutero \, phosphoric \, acid} \end{array}$$

$$SO_3 + D_2O \longrightarrow D_2SO_4$$
Deutero sulphuric acid

Action with metallic carbides, phosphides, nitrides, etc. : Like H2O heavy water (d) reacts with carbides, phosphide nitrides, arsenides, etc. to form corresponding deutero compounds.

$$CaC_2 + 2D_2O \longrightarrow Ca(OD)_2 + C_2D$$
Deutero algorithme

$$Al_4C_3 + 12D_2O \longrightarrow 4Al(OD)_3$$
Deuts o methane

$$Mg_2N_2 + 6D_2O \longrightarrow 3Mg(OD)_2$$
 $2ND_3$

$$Mg_3N_2 + 6D_2O \longrightarrow 3Mg(OD)_2 2ND_3$$
Deutero ammon

$$AIN + 3D_2O \longrightarrow AI(OD)_3 + ND_3$$
Deutero ammonia

$$Ca_3P_2 + 6D_2O \longrightarrow 3Ca(OD)_2 + 2PD_3$$
Deutero phosphine

$$Na_3As + 3D_2O \longrightarrow 3NaOD + AsD_3$$
Deutero ar sir

Electrolysis. A solution of heavy water containing Na2CO3 when electrolysed evolve heavy hydrogen at (e) cathode

$$2D_2O \xrightarrow{\text{Electrolysis}} 2D_2 + O_2$$
(Cathode) (Anode

Uses: As a neutron moderator: Fission in uranium-235 is brought by slow speed neutrons. The substances which are used for slowing down the speed of neutrons are called moderators. Heavy water is used for this purpose in nuclear reactors.

H₂O₂ (Hydrogen Peroxide)

Laboratory method:

In laboratory, H_2O_2 is prepared by adding calculated amounts of sodium peroxide to ice cod dilute 20%, solution of H_2SO_4 .

$$Na_2O_2 + H_2SO_4 \longrightarrow Na_2SO_4 + H_2O_2$$

By the action of sulphuric acid or phosphoric acid on hydrated barium peroxide BaO₂.8H₂O (Meros process)

(a)
$$BaO_2.8H_2O + H_2SO_4 \longrightarrow BaSO_4 \downarrow +H_2O_2 + 8H_2O$$

Anhydrous barium peroxide does not react readity with sulphuric acid (because a coating of insoluble barium sulphate is formed on its surface which stops further action of the acid). Therefore, hydrated barium peroxide, $BaO_2.8H_2O$ must be used.

(b)
$$3\text{BaO}_2 + 2\text{H}_3\text{PO}_4 \longrightarrow \text{Ba}_3(\text{PO}_4)_2 + 3\text{H}_2\text{O}$$

 $\text{Ba}_3(\text{PO}_4)_2 + 3\text{H}_2\text{SO}_4 \longrightarrow 3\text{BaSO}_4 + 2\text{H}_3\text{PO}_4$

Phosphoric acid is preferred over H_2SO_4 because soluble impurities like barium persulphate $(BaO_2.8H_2O_4.8H_2O_4)$ tends to decompose H_2O_2 while H_3PO_4 acts as preservative (negative catalyst) for H_2O_2 .

Industrial Method

(i) Auto oxidation of 2 butyl anthraquinol (cyclic process):

$$C_{4}H_{9}$$

$$OH$$

$$C_{4}H_{9}$$

$$OH$$

$$C_{4}H_{2}$$

(ii) Oxidation of isopropyl alcohol:

(iii) Electrolytic Process: Used 50% HSO in electrolytic cell using Pt as anode graphite as cathode.

$$2H_{2}SO_{4} \longrightarrow 2H_{1} + 2HSO_{4}^{\Theta}$$
At Cathode $2H^{+} + 2e^{-} \longrightarrow H_{2}S_{2}O_{8} + 2e^{-}$

At Anode $2HSO_{4}^{-} \longrightarrow H_{2}S_{2}O_{8} + 2e^{-}$

Peroxo disulphhuric acid

$$H_{2}S_{2}O_{8} + H_{2}O_{2} \longrightarrow H_{2}SO_{4} + H_{2}SO_{8}$$

$$H_{3}SO_{4} \longrightarrow H_{2}SO_{4} + H_{3}SO_{4} + H_{3}SO_{4} + H_{3}SO_{4} + H_{4}SO_{5}$$

Physical property

- (i) Pure H_2O_2 is colour less, odourless liquid and impure with bluish layer.
- (ii) It has more Hydrogen bonding then H_2O , So order of boiling point. $H_2O_2 > D_2O > H_2O$
- (iii) It is soluble in H₂O, alcohol and ether
- (iv) It has bitter test and harmful for skin
- (v) It is a dibasic weak acid
- (yi) It has a oxidising as well as reducing property
- (vii) $H_2^2O_2$ easily decompose in presence of light and temperature, So H_2O_2 always kep in dark bottles and kept at cool places.
- (viii) 30% solution of H_2O_2 is called **Perhydrol**.

Chemical property:

1. Stabily: It is unstable in nature decompose on standing and heating. It is an example of auto oxidation-

reduction Reaction; $H_2O_2 \rightarrow H_2O + O$

Note: Nascent oxygen working as colourless agent

Decomposition of H_2O_2 ; $2H_2O_2 \rightleftharpoons 2H_2O + O_2$

Note: This is retarded by R-OH, acetenilide, glycerol

2. Oxidizing nature: H₂O₂ is a powerful oxidant in acidic as well as in alkaline medium.

 $H_{2}O_{2} + 2H^{+} + 2e \xrightarrow{Fast} 2H_{2}O \qquad E_{RD}^{0} = 1.77V$ (In acid)

 $H_2O + HO_2^- + 2e \xrightarrow{Slow} 3HO^- \qquad E_{RP}^0 = 0.87V$ (In alkali)

Thuse H_2O_2 is more powerful oxidant in acidic medium. The simple interpretation of O_2O_2 as oxidant can be shown by the equation.

$$H_2O_2 \longrightarrow H_2O + O$$

Following are some important examples of oxidant action of H_2O_2 :

 $\mathrm{H_2O_2}$ oxidises black lead sulphide (PbS) to white lead sulphate (PbSO). This reaction is used in restoring the white colour of old paintings which have blackened due to the formation of lead sulphide by the action of H_0S present in air.

> $4[H_2O_2 \longrightarrow H_2O + O]$ $PbS + 4O \longrightarrow PbSO_{4}$

$$PbS + 4H_2O_2 \longrightarrow PbSO_4 + 4H_2O_4$$

 H_9O_9 oxidises sulphites into sulphates. (b)

> $H_2O_2 \longrightarrow H_2O + O$ $Na_2SO_3 + O \longrightarrow Na_2SO_4$

$$Na_2SO_3 + H_2O_2$$
 $Na_2SO_4 + H_2O_3$

H₂O₂ oxidises nitrites to intrates (c)

$$Na_2NO_2 + H_2O_2 \longrightarrow Na_2NO_3 + H_2O$$

H,O, oxidises N,S into sulphur. (d)

 $H_2O_2 \longrightarrow H_2O + O$ $H_2S + O \longrightarrow H_2O + S$

$$H_2S + O \longrightarrow H_2O + S$$

$$H_2S + H_2O_2 \longrightarrow 2H_2O + S$$

Reducing nature: It can also act as a reducing agent towards powerful oxidising agents. 3.

 $H_2O_2 \longrightarrow 2H^+ + O_2^- + 2e^-$

It reduces Ag₂O to silver.

 $Ag_2O + H_2O_2 \longrightarrow 2Ag + H_2O + O_2$

It reduces ozone to oxygen.

 $H_{9}O_{9} + O_{3} \longrightarrow H_{9}O + 2O_{9}$

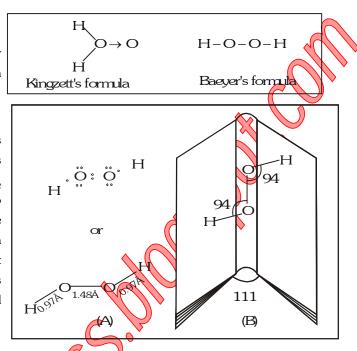
Structure of Hydrogen Peroxide

The vapour density as determined by Victor

Meyermethod at 90 C is 17. Hence, the molecular mass of $\rm H_2O_2$ is 34. Two formulae have been suggested for hydrogen peroxide.

The calculated value of the single bond O-O

distance is 1.48Å and X-ray measurements shows what in hydrogen peroxide, O–O bond distance is 1.46 + 0.03Å. The value of dipole moment of $\rm H_2O_2$ is 2.1 D. This suggests that all the four atoms do not lie in the same plane. The molecule can be pictured as lying on the spine of a book open to an angle of 111 . The hydrogen atoms are present one on each cover and H–O bonds making angles of 94 with the O–O bond as shown in fig. The bond distance between O–H is 0.97Å.



Precautions:

(i) H_2O_2 can not be stored in simple glass bottles since rough surface of glass [alkali oxides present in it] excited by light and decomposed H_2O_2 .

So the H2O2 usually stored in coloured, paraffin wax coated, plastic bottle.

(ii) Always with H_2O_2 add small quantity of inhibitor or negative catalyst to stay decomposition of H_2O_2 .

like H_3PO_4 , acetanilides etc.

	Uses		
(i)	Bleaching agent	(ii)	Hair dying
(iii)	$H_2O_2 + N_2H_4$ as Rocket propellent	(iv)	$\boldsymbol{H}_{\!\scriptscriptstyle 2}\boldsymbol{O}_{\!\scriptscriptstyle 2}$ as oxidant and reductant
(v)	Antiseptic		

DIHYDROGEN AS A FUEL

Dihydrogen releases large quantities of heat on combustion. The data on energy released by combustion of fuels like dihydrogen, methane, LPG etc. are compared in terms of the same amounts in mole, mass and volume, are shown in Table

THE ENERGY RELEASED BY COMBUSTION OF VARIOUS FUELS IN MOLES, MASS AND VOLUME

Energy released on combustion in kJ state)	Dihydrogen (in gaseous state)	Dihydrogen (in liquid)	LPG	CH ₄ gas	Octane (in liquid state)
per mole	286	285	2220	880	5511
per gram	143	142	50	53	47
per litre	12	9968	25590	35	34005

From this table it is clear that on a mass for mass basis dihydrogen can release more energy than petrol (about three times). Moreover, pollutants in combustion of dihydrogen will be less than petrol. The only pollutants will be the oxides of dinitrogen (due to the presence of dinitrogen as impurity with dihydrogen). This, of course, can be minimised by injecting a small amount of water into the cylinder to lower the temperature so that the reaction between dinitrogen and dioxygen may not take place. However, the mass of the containers in which dihydrogen will be kept must be taken into consideration. A cylinder of compressed dihydrogen weighs about 30 times as much as a tank of petrol containing the same amount of energy. Also, dihydrogen gas is converted into liquid state by cooling to 20K. This would require expensive insulated tanks. Tanks of metal alloy like NaNi₅, Ti-TiH, Mg-MgH₂ etc. are in use for storage of dihydrogen in small quantities. These limitations have prompted researchers to search for alternative techniques to use dihydrogen in an efficient way.

In this view **Hydrogen Economy** is an alternative. The basic principle of hydrogen economy is the transportation and storage of energy in the form of liquid or gaseous dihydrogen. Advantage of hydrogen economy is that energy is transmitted in the form of dihydrogen and not as electric power. It is for the first time in the history of India that a pilot project using dihydrogen as fuel was launched in October 2005 for running automobiles. Initially 5% dihydrogen has been mixed in CNG for use in four-wheeler vehicles. The percentage of dihydrogen would be gradually increased to reach the optimum level.

Nowadays, it is also used in fuel cells for generation of electric power. It is expected that economically viable and safe sources of dihydrogen will be identified in the years to come, for its usage as a common source of energy.



MISCELLANEOUS SOLVED EXAMPLES

- 1. H_2O_2 and heavy water was discovered by respectively:-
 - (1) Thenard, Urey
- (2) Urey, Rutherford
- (3) Aston, Urey
- (4) Aston, Chadwick

Sol. (1)

> $\mathrm{H_2O_2}$ was discovered by Thenard where was Urey discovered heavy water.

 $4D_2O + 3Fe \xrightarrow{red-hot} Fe_3O_4 + gas.$ 2.

The gas produced in the above reaction is :-

- (1) O_2
- (2) H_2
- (3) D_2
- (4) None

Sol. (3)

The gas produced in the reaction is deuterium.

- 3. Which of the following compound affects mercury
 - (1) D₂O
- (2) $H_{2}O$
- (3) O_3
- (4) dil. HCl

(3) Sol.

> Ozone wipes out meniscus of mercury due to the formation of Hg₂O as a result of [O₃ - Hg] reaction

- 4. An inorganic substance liberates oxygen on heating and turns an acidic solution of KI brown and reduces acidified $KMnO_4$ solution. The substance is :-
 - (1) HgO
- (2) H_2O_2
- (3) KNO₃
- (4) $Pb(NO_3)_2$

Sol. (2)

$$2H_2O_2 \xrightarrow{\text{Heat}} 2H_2O + O_2$$
;

$$2KI + H_2O_2 \xrightarrow{Acidic} 2KOH + I_2$$

Acidified KMnO₄ acts as an oxidising agent when it is decolourised, $H_2O_2 + O_2 + O_2$

- 5. In which of the following reactions does hydrogen act as an oxidising agent
 - (1) $H_2 + F_2 \longrightarrow$
- (2) $H_2 + SiCl_4 \longrightarrow$ (4) $CuO + H_2 \longrightarrow$
- (3) Na + H

Sol. (3)

> The reaction in which hydrogen accepts the electron or its exidation number changes from 0 , it acts as an oxidising agent.

 $a + H_2 \longrightarrow 2Na + H^-$

- When H2O2 is added to ice cold solution of aciditied 6. potassium dichromate in ether and the contents are shaken and allowed to stand :
 - (1) A blue colour is obtained in ether due to formation of $Cr_2(SO_4)_3$
 - (2) A blue colour is obtained in either due to formation of CrO₅
 - (3) A blue colour is obtained in other due to formation of CrO₃
 - (4) Chromyl chloride is formed
- Sol. (2)

$$\operatorname{Cr_2O_7^{2-}} + 2\operatorname{H}^{\dagger} + 4\operatorname{H_2O_2} \rightarrow 2\operatorname{CrO_5}_{\text{Blue colour}} + 5\operatorname{H_2O_2}$$

- 7. When a substance A reacts with water, it produces a combustible gas B and a solution of substance C In water. D reacts with this solution of C and produces the same gas B on warming. D can also produce gas B on reaction with dilute H_2SO_4 . A imparts a deep golden yellow colour to smokeless flame. A, B, C and D respectively are :-
 - (1) Na, H₂, NaOH, Zn
 - (2) K, H₂, KOH, Al
 - (3) Ca, H₂, Ca(OH)₂, Sn
 - (4) CaC₂, C₂H₂, Ca(OH)₂, Fe
- Sol. (1)

$$Zn + H_2SO_4 \longrightarrow ZnSO_4 + H_2$$
(B) (B)