• The s-block elements of the Periodic Table are those in which the last electron enters the outermost s-orbital. As the s-orbital can accommodate only two electrons, two groups (1 & 2) belong to the s-block of the Periodic Table.

Group 1 of the Periodic Table consists of the elements:

 lithium, sodium, potassium, rubidium, caesium and francium. They are collectively known as the alkali metals.

- These are so called because they form hydroxides on reaction with water which are strongly alkaline in nature.
- The elements of Group 2 include beryllium, magnesium, calcium, strontium, barium and radium. These elements with the exception of beryllium are commonly known as the alkaline earth metals. These are so called because their oxides and hydroxides are alkaline in nature and these metal oxides are found in the earth's crust*.

- Among the alkali metals sodium and potassium are abundant and lithium, rubidium and caesium have much lower abundances
- Francium is highly radioactive; its longest-lived isotope 223Fr has a half-life of only 21 minutes.
- Of the alkaline earth metals calcium and magnesium ranks fifth and sixth in abundance respectively in the earths crust.
- Strontium and barium have much lower abundances. Beryllium is rare and radium is the rarest of all comprising only 10⁻¹⁰ per cent of igneous rocks†

- The general electronic configuration of s-block elements is [**noble gas**]**ns**¹ for alkali metals and [**noble gas**] **ns**² for alkaline earth metals.
- Lithium and beryllium, the first elements of Group 1 and Group 2 respectively exhibit some properties which are different from those of the other members of the respective group.
- In these anomalous properties they resemble the second element of the following group.

 Thus, lithium shows similarities to magnesium and beryllium to aluminium in many of their properties. This type of diagonal similarity is commonly referred to as diagonal relationship in the periodic table.

• The diagonal relationship is due to the similarity in ionic sizes and /or

charge/radius ratio of the elements.

 Monovalent sodium and potassium ions and divalent magnesium and calcium ions are found in large proportions in biological fluids.

• These ions perform important biological functions such as maintenance of ion balance and nerve impulse conduction.

GROUP 1 ELEMENTS: ALKALI METALS

- The alkali metals show regular trends in their physical and chemical properties with the increasing atomic number.
- The atomic, physical and chemical properties of alkali metals are discussed below.

Electronic Configuration

Element	Symbol	Electronic configuration		
Lithium	Li	1s ² 2s ¹		
Sodium	Na	$1s^22s^22p^63s^1$		
Potassium	K	$1s^22s^22p^63s^23p^64s^1$		
Rubidium	Rb	$1s^22s^22p^63s^23p^64s^23d^{10}4p^65s^1$		
Caesium	Cs	$1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^{10} 4p^6 4d^{10} 5s^2 5p^6 6s^1$		
		or [Xe] 6s ¹		
Francium	Fr	[Rn]7s ¹		

Electronic Configuration

- All the alkali metals have one valence electron, ns¹ outside the noble gas core.
- The loosely held s-electron in the outermost valence shell of these elements makes them the most electropositive metals. They readily lose electron to give monovalent M⁺ ions. Hence they are never found in Free State in nature.

Atomic and Ionic Radii

- The alkali metal atoms have the largest sizes in a particular period of the periodic table. With increase in atomic number, the atom becomes larger.
- The monovalent ions (M⁺) are smaller than the parent atom.
- The atomic and ionic radii of alkali metals increase on moving down the group i.e., they increase in size while going from Li to Cs.

Ionization Enthalpy

 The ionization enthalpies of the alkali metals are considerably low and decrease down the group from Li to Cs.

 This is because the effect of increasing size outweighs the increasing nuclear charge, and the outermost electron is very well screened from the nuclear charge.

Hydration Enthalpy

• The hydration enthalpies of alkali metal ions decrease with increase in ionic sizes.

• $Li^+> Na^+> K^+> Rb^+> Cs^+$

• Li⁺ has maximum degree of hydration and for this reason lithium salts are mostly hydrated, e.g., LiCl·2H₂O

Physical Properties

- All the alkali metals are silvery white, soft and light metals.
- Because of the large size, these elements have low density which increases down the group from Li to Cs. However, potassium is lighter than sodium.
- The melting and boiling points of the alkali metals are low indicating weak metallic bonding due to the presence of only a single valence electron in them.

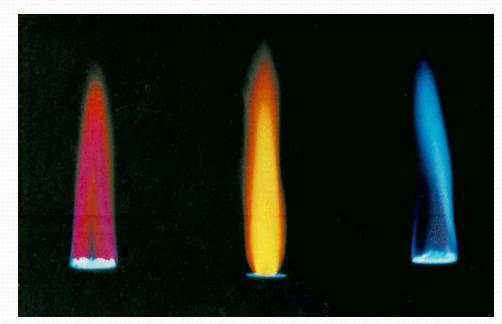
Physical Properties

- The alkali metals and their salts impart characteristic colour to an oxidizing flame.
- This is because the heat from the flame excites the outermost orbital electron to a higher energy level. When the excited electron comes back to the ground state, there is emission of radiation in the visible region as given below:

Flame tests







Alkali metals can therefore, be detected by the respective flame tests and can be determined by flame photometry or atomic absorption spectroscopy.

• These elements when irradiated with light, the light energy absorbed may be sufficient to make an atom lose electron. This property makes caesium and potassium useful as electrodes in photoelectric cells.

Table 10.1 Atomic and Physical Properties of the Alkali Metals

Property	Lithium	Sodium	Potassium	Rubidium	Caesium	Francium		
	Li	Na	К	Rb	Cs	Fr		
Atomic number	3	11	19	37	55	87		
Atomic mass (g moΓ¹)	6.94	22.99	39.10	85.47	132.91	(223)		
Electronic configuration	[He] 2s ¹	[Ne] 3s ¹	[Ar] 4s ¹	[Kr] 5s ¹	[Xe] 6s ¹	[Rn] 7s ¹		
lonization enthalpy / kJ mol ⁻¹	520	496	419	403	376	~375		
Hydration enthalpy/kJ mol ⁻¹	-506	-406	-330	-310	-276	_		
Metallic radius / pm	152	186	227	248	265	_		
Ionic radius M ⁺ / pm	76	102	138	152	167	(180)		
m.p. / K	454	371	336	312	302	-		
b.p / K	1615	1156	1032	961	944	-		
Density / g cm ⁻³	0.53	0.97	0.86	1.53	1.90	-		
Standard potentials E ⁰ / V for (M ⁺ / M)	-3.04	-2.714	-2.925	-2.930	-2.927	-		
Occurrence in lithosphere [†]	18*	2.27**	1.84**	78-12*	2-6*	~ 10 ⁻¹⁸ *		

^{*}ppm (part per million), ** percentage by weight; † Lithosphere: The Earth's outer layer: its crust and part of the upper mantle

Chemical Properties

 The alkali metals are highly reactive due to their <u>large size and low</u> <u>ionization enthalpy</u>.

 The reactivity of these metals <u>increases</u> down the group.

(i) Reactivity towards air:

• The alkali metals tarnish in dry air due to the formation of their oxides which in turn react with moisture to form hydroxides.

They burn vigorously in oxygen forming oxides.

• Lithium forms monoxide, sodium forms peroxide, the other metals form superoxides. The superoxide O_2^- ion is stable only in the presence of large cations such as K, Rb, Cs.

•
$$4\text{Li} + O_2 \rightarrow 2\text{Li}_2O$$
 (oxide)

•
$$2Na+O_2 \rightarrow Na_2O_2$$
 (peroxide)

•
$$M+O_2 \rightarrow MO_2$$
 (superoxide)
(M = K, Rb, Cs)

- In all these oxides the oxidation state of the alkali metal is +1.
- Lithium shows exceptional behaviour in reacting directly with nitrogen of air to form the nitride, Li₃N as well

Note = (oxide)
$$O^{2-}$$
 (peroxide) O^{-1} (superoxide)

Because of their high reactivity towards air and water, they are normally kept in kerosene oil.

(ii) Reactivity towards water:

The alkali metals react with water to form hydroxide and dihydrogen.

$$2M + 2H_2O \rightarrow 2M^+ + 2OH^- + H_2$$

(M = an alkali metal)

- although lithium has most negative E° value its reaction with water is less vigorous than that of sodium which has the least negative E° value among the alkali metals.
- This behaviour of lithium is attributed to its small size and very high hydration energy.
- Other metals of the group react explosively with water.
- They also react with proton donors such as alcohol, gaseous ammonia and alkynes.

(iii) Reactivity towards dihydrogen:

• The alkali metals react with dihydrogen at about 673K (lithium at 1073K) to form hydrides.

 All the alkali metal hydrides are ionic solids with high melting points.

$${\color{red} \bullet}$$
 2M + H₂ \rightarrow 2M⁺+H⁻

(iv) Reactivity towards halogens

- The alkali metals readily react vigorously with halogens to form ionic halides, M+X- However, lithium halides are somewhat covalent. It is because of the high polarisation capability of lithium ion (The distortion of electron cloud of the anion by the cation is called polarisation).
- The Li⁺ ion is very small in size and has high tendency to distort electron cloud around the negative halide ion. Since anion with large size can be easily distorted, among halides, lithium iodide is the most covalent in nature.

Uses:

- Lithium metal is used to make useful alloys, for example with lead to make 'white metal' bearings for motor engines, with aluminium to make aircraft parts, and with magnesium to make armour plates.
- Lithium is used in thermonuclear reactions.
- Lithium is also used to make electrochemical cells.
- Sodium is used to make a Na/Pb alloy needed to make PbEt₄ and PbMe₄.
- These organolead compounds were earlier used as anti-knock additives to petrol, but nowadays vehicles use lead-free petrol

Uses: ----contd

- Liquid sodium metal is used as a coolant in fast breeder nuclear reactors.
- Potassium has a vital role in biological systems.
- Potassium chloride is used as a fertilizer.
- Potassium hydroxide is used in the manufacture of soft soap.
- Potassium hydroxide is also used as an excellent absorbent of carbon dioxide.
- Caesium is used in devising photoelectric

5-Mar-20 cells.

WHY LITHIUM SHOWS ANOMALOUS PROPERTIES

- REASON FOR ANOMALOUS BEHAVIOUR
- The anomalous behavior of lithium is due to the:
- exceptionally small size of its atom and ion, and
- high polarizing power (i.e., charge/ radius ratio)

As a result, there is increased covalent character of lithium compounds which is responsible for their solubility in organic solvents.

ANOMALOUS PROPERTIES OF LITHIUM

- (i) Lithium is much harder. Its m.p. and b.p. are higher than the other alkali metals.
- (ii) Lithium is least reactive but the strongest reducing agent among all the alkali metals.On combustion in air it forms mainly monoxide, Li₂O and the nitride, Li₃N unlike other alkali metals.
- (iii) LiCl is deliquescent and crystallises as a hydrate, LiCl.2H₂O whereas other alkali metal chlorides do not form hydrates.
- (iv) Lithium hydrogencarbonate is not obtained in the solid form while all other elements form solid hydrogencarbonates.

- (v) Lithium unlike other alkali metals forms no ethynide on reaction with ethyne.
- (vi) Lithium nitrate when heated gives lithium oxide, Li₂O, whereas other alkali metal nitrates decompose to give the corresponding nitrite.

$$4LiNO3 \rightarrow 2Li2O + 4NO2 + O2$$

$$2 NaNO3 \rightarrow 2NaNO2 + O2$$

 (vii) LiF and Li₂O are comparatively much less soluble in water than the corresponding compounds of other alkali metals.

Points of Similarities between Lithium and Magnesium / Diagonal Relationship

 The similarity between lithium and magnesium is particularly striking and arises because of their similar sizes

```
Atomic radii, Li = 152 pm, Mg = 160 pm; ionic radii : Li<sup>+</sup> = 76 pm, Mg<sup>2+</sup> = 72 pm.
```

The main points of similarity are

- (i) Both lithium and magnesium are harder and lighter than other elements in the respective groups.
- (ii) Lithium and magnesium react slowly with water. Their oxides and hydroxides are much less soluble and their hydroxides decompose on heating. Both form a nitride, Li₃N and Mg₃N₂, by direct combination with nitrogen.
- (iii) The oxides, Li₂O and MgO do not combine with excess oxygen to give any superoxide.

(iv) The carbonates of lithium and magnesium decompose easily on heating to form the oxides and CO₂. Solid hydrogen carbonates are not formed by lithium and magnesium.

• (v) Both LiCl and MgCl₂ are soluble in ethanol.

• (vi) Both LiCl and MgCl₂ are deliquescent and crystallize from aqueous solution as hydrates, LiCl·2H₂O and MgCl₂·8H₂O.

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SOME IMPORTANT COMPOUNDS OF SODIUM

- Sodium carbonate,

 Na₂CO₃·10H₂O
- Sodium hydroxide,NaOH
- Sodium chloride and <u>NaCl</u>
- Sodium bicarbonate
 NaHCO₃

Sodium Carbonate (Washing Soda),

$Na_2CO_3 \cdot 10H_2O$

- Sodium carbonate is generally prepared by Solvay Process.
- In this process, advantage is taken of the low solubility of sodium hydrogencarbonate whereby it gets precipitated in the reaction of sodium chloride with ammonium hydrogencarbonate.
- The latter is prepared by passing CO₂ to a concentrated solution of sodium chloride saturated with ammonia, where ammonium carbonate followed by ammonium hydrogencarbonate are formed.

 The equations for the complete process may be written as :

$$\begin{split} 2\,\mathrm{NH_3} + \mathrm{H_2O} + \mathrm{CO_2} &\to (\mathrm{NH_4})_2\,\mathrm{CO_3} \\ (\mathrm{NH_4})_2\,\mathrm{CO_3} + \mathrm{H_2O} + \mathrm{CO_2} &\to 2\,\mathrm{NH_4HCO_3} \\ \mathrm{NH_4HCO_3} + \mathrm{NaCl} &\to \mathrm{NH_4Cl} + \mathrm{NaHCO_3} \end{split}$$

Sodium hydrogencarbonate crystal separates.
 These are heated to give sodium carbonate.

$$2NaHCO_3 \rightarrow Na_2CO_3 + CO_2 + H_2O$$

In this process NH₃ is recovered when the solution containing NH₄Cl is treated with Ca(OH)₂. Calcium chloride is obtained as a byproduct.

$$2NH_4Cl + Ca(OH)_2 \rightarrow 2NH_3 + CaCl_2 + H_2O$$

Solvay process cannot be extended to the manufacture of potassium carbonate because potassium hydrogencarbonate is too soluble to be precipitated by the addition of ammonium hydrogencarbonate to a saturated solution of potassium chloride

Properties:

• Sodium carbonate is a white crystalline solid which exists as a decahydrate,

 $Na_2CO_3 \cdot 10H_2O.$

This is also readily soludecated anydrate to form moments and anydrous called soda

da. It is ating, the crystallization 73K, the etely nite powder

 Carbonate part of sodium carbonate gets hydrolysed by water to form an alkaline solution.

$$CO_3^{2-} + H_2O \rightarrow HCO_3^- + OH^-$$

Uses:

- (i) It is used in water softening, laundering and cleaning.
- (ii) It is used in the manufacture of glass, soap, borax and caustic soda.
- (iii) It is used in paper, paints and textile industries.
- (iv) It is an important laboratory reagent both in qualitative and quantitative analysis.

Sodium Chloride, NaCl



Calcium chloride, CaCl₂, and magnesium chloride, MgCl₂ are impurities because they are deliquescent (absorb moisture easily from the atmosphere).

 To obtain pure sodium chloride, the crude salt is disso amount of nsoluble water and impurities en saturated with hydro rystals of pure sodium ch Calcium and magnesium cmoriue, being more soluble than sodium chloride, remain in solution.

Properties:

- Sodium chloride melts at 1081K.
- It has a solubility of 36.0 g in 100 g of water at 273 K.
- The solubility does not increase appreciably with increase in temperature.

Uses:

- (i) It is used as a common salt or table salt for domestic purpose.
- (ii) It is used for the preparation of Na₂O₂, Na₂OH and Na₂CO₃.

Sodium Hydroxide (Caustic Soda), NaOH

- Sodium hydroxide is generally prepared commercially by the electrolysis of sodium chloride in Castner-Kellner cell.
- A brine solution is electrolysed using a mercury cathode and a carbon anode.
 Sodium metal discharged at the cathode combines with mercury to form sodium amalgam.
- •••Chlorine gas is evolved at the anode.

Cell reactions

Cathode:
$$Na^+ + e^- \xrightarrow{Hg} Na - amalgam$$

Anode :
$$Cl^- \rightarrow \frac{1}{2}Cl_2 + e^-$$

The amalgam is treated with water to give sodium hydroxide and hydrogen gas.

2Na-amalgam + 2H₂O 2NaOH+ 2Hg +H₂

Properties:

- Sodium hydroxide is a white, translucent solid.
- It melts at 591 K.
- It is readily soluble in water to give a strong alkaline solu

de are

- Crystals of s deliquescent
- The sodium tion at the surface reacts with the CO₂ in the atmosphere to form Na₂CO₂.

Uses:

- It is used in the manufacture of soap,paper, artificial silk and a number of chemicals,
- in petroleum refining,
- in the purification of bauxite,
- in the textile industries for mercerizing cotton fabrics,
- for the preparation of pure fats and oils, and
- ⁵ as a laboratory reagent.

Sodium Hydrogencarbonate (BakingSoda), NaHCO₃

Sodium hydrogencarbonate is known

as bak on hea carbon or pas fluffy)



mposes of n cakes ght and

- Sodium hydrogencarbonate is made by saturating a solution of sodium carbonate with carbon dioxide.
- The white crystalline powder of sodium hydrogencarbonate, being less soluble, gets separated out.

$$Na_2CO_3+H_2O+CO_2\rightarrow 2NaHCO_3$$

- Sodium hydrogencarbonate is a mild antiseptic for skin infections.
- It is used in fire extinguishers.

BIOLOGICAL IMPORTANCE OF

SODIUM AND POTASSIUM

• A typical 70 kg man contains about 90 g of Na and 170 g of K compared with only 5 g of iron and 0.06 g of copper.

 Sodium ions are found primarily on the outside of cells, being located in blood plasma and in the interstitial fluid which surrounds the cells.

SODIUM AND POTASSIUM

- These ions participate in the transmission of nerve signals, in regulating the flow of water across cell membranes and in the transport of sugars and amino acids into cells.
- Sodium and potassium, although so similar chemically, differ quantitatively in their ability to penetrate cell membranes, in their transport mechanisms and in their efficiency to activate enzymes.

SODIUM AND POTASSIUM

 Thus, potassium ions are the most abundant cations within cell fluids, where they activate many enzymes, participate in the oxidation of glucose to produce ATP and, with sodium, are responsible for the transmission of nerve signals.

 There is a very considerable variation in the concentration of sodium and potassium ions found on the opposite
 sides of cell membranes.

SODIUM AND POTASSIUM

- As a typical example, in blood plasma, sodium is present to the extent of 143 mmolL⁻¹, whereas the potassium level is only 5 mmolL⁻¹ within the red blood cells.
- These concentrations change to 10 mmolL⁻¹ (Na⁺) and 105 mmolL⁻¹ (K⁺). These ionic gradients demonstrate that a discriminatory mechanism, called the sodium-potassium pump, operates across the cell membranes which consumes more than one-third of the ATP used by a resting animal and about 15 kg per 24 hour in a resting human.
- 5-MMOTE:- mmolL-1 = mili moles per litre)

GROUP 2 ELEMENTS:

ALKALINE EARTH METALS

• The group 2 elements comprise beryllium, magnesium, calcium, strontium, barium and radium. They follow alkali metals in the periodic table. These (except beryllium) are known as alkaline earth metals. The first element beryllium differs from the rest of the members and shows diagonal relationship to aluminium.

Electronic configuration

- These elements have two electrons in the s -orbital of the valence shell.
- •Their general electronic configuration may be represented as [noble gas] ns².
- Like alkali metals, the compounds of these elements are also predominantly ionic.

Electronic Configuration

Element	Symbol	Electronic configuration
Beryllium	Be	1s ² 2s ²
Magnesium	Mg	$1s^22s^22p^63s^2$
Calcium	Ca	$1s^22s^22p^63s^23p^64s^2$
Strontium	Sr	$1s^22s^22p^63s^23p^64s^23d^{10}4p^65s^2$
Barium	Ba	$1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^{10} 4p^6 4d^{10} 5s^2 5p^6 6s^2$
		or [Xe] 6s ²
Radium	Ra	[Rn]7s ²

Element	Symbol	Electronic configuration
Beryllium	Be	$1 s^2 2 s^2$
Magnestum	Mg	$1 s^2 2 s^2 2 p^6 3 s^2$
Calcium	Ca	$1s^22s^22p^63s^23p^64s^2$
Strontium	Sr	1 s ² 2 s ² 2 p ⁶ 3 s ² 3 p ⁶ 3 d ¹⁰ 4 s ² 4 p ⁶ 5 s ²
Bartum	Ba	1 s ² 2 s ² 2 p ⁶ 3 s ² 3 p ⁶ 3 d ¹⁰ 4 s ² 4 p ⁶ 4 d ¹⁰ 5 s ² 5 p ⁶ 6 s ² or [Xe]6 s ²
Radium	Ra	[Rn]7s ²

Atomic and Ionic Radii

- The atomic and ionic radii of the alkaline earth metals are smaller than those of the corresponding alkali metals in the same periods. This is due to the increased nuclear charge in these elements.
- Within the group, the atomic and ionic radii increase with increase in atomic number.

Ionization Enthalpies

- The alkaline earth metals have low ionization enthalpies due to fairly large size of the atoms.
 Since the atomic size increases down the group, their ionization enthalpy decreases
- The first ionisation enthalpies of the alkaline earth metals are higher than those of the corresponding Group 1 metals. This is due to their small size as compared to the corresponding alkali metals. It is interesting to note that the second ionisation enthalpies of the alkaline earth metals are smaller than those of the corresponding alkali metals.

Hydration Enthalpies
Like alkali metal ions, the hydration enthalpies of alkaline earth metal ions decrease with increase in ionic size down the group.

$$Be^{2+} > Mg^{2+} > Ca^{2+} > Sr^{2+} > Ba^{2+}$$

 The hydration enthalpies of alkaline earth metal ions are larger than those of alkali metal ions. Thus, compounds of alkaline earth metals are more extensively hydrated than those of alkali metals, e.g., MgCl, and CaCl, exist as MgCl, 6H,O and CaCl, 6H,O while NaCl and KCl do not

5-Marform such hydrates.

Physical Properties

- The alkaline earth metals, in general, are silvery white, lustrous and relatively soft but harder than the alkali metals.
- Beryllium and magnesium appear to be somewhat grayish.
- The melting and boiling points of these metals are higher than the corresponding alkali metals due to smaller sizes. The trend is, however, not systematic

Physical Properties contd

- Because of the low ionization enthalpies, they are strongly electropositive in nature.
- The electropositive character increases down the group from Be to Ba.
- Calcium, strontium and barium impart characteristic brick red, crimson and apple green colours respectively to the flame.

Physical Properties contd

- The electrons in beryllium and magnesium are too strongly bound to get excited by flame. Hence, these elements do not impart any colour to the flame.
- The flame test for Ca, Sr and Ba is helpful in their detection in qualitative analysis and estimation by flame photometry.
- The alkaline earth metals like those of alkali metals have high electrical and thermal conductivities which are typical characteristics of metals.

Chemical Properties

 The alkaline earth metals are less reactive than the alkali metals. The reactivity of these elements increases on going down the group.

(i) Reactivity towards air and water:

- Beryllium and magnesium are kinetically inert to oxygen and water because of the formation of an oxide film on their surface.
- However, powdered beryllium burns brilliantly on ignition in air to give BeO 5-Marand Be₃N₂.

Chemical Properties

- Magnesium is more electropositive and burns with dazzling brilliance in air to give MgO and Mg₃N₂.
- Calcium, strontium and barium are readily attacked by air to form the oxide and nitride.
- They also react with water with increasing vigour even in cold to form hydroxides.

(ii) Reactivity towards the halogens:

 All the alkaline earth metals combine with halogen at elevated temperatures forming their halides

$$M+X_2 \rightarrow MX_2$$
 (X=F,Cl,Br,l)

Thermal decomposition of (NH₄)₂BeF₄ is the best route for the preparation of BeF₂, and BeCl₂ is conveniently made from the oxide.

(iii) Reactivity towards hydrogen:

All the elements except beryllium combine with hydrogen upon heating to form their hydrides, MH₂.BeH₂, however, can be prepared by the reaction of BeCl₂ with LiAlH₄.

$$2\text{BeCl}_2 + \text{LiAlH}_4 \rightarrow 2\text{BeH}_2 + \text{LiCl} + \text{AlCl}_3$$

(iv) Reactivity towards acids:

• The alkaline earth metals readily react with acids liberating dihydrogen.

$$M + 2HCl \rightarrow 2MCl_2 + H_2$$

- Beryllium is used in the manufacture of alloys.
- Copper-beryllium alloys are used in the preparation of high strength springs.
- Metallic beryllium is used for making windows of X-ray tubes.
- Magnesium-aluminium alloys being light in mass are used in air-craft construction
- Radium salts are used in radiotherapy, for example, in the treatment of cancer.

Uses:

- Magnesium (powder and ribbon) is used in flash powders and bulbs, incendiary bombs and signals.
- A suspension of magnesium hydroxide in water (called milk of magnesia) is used as antacid in medicine.
- Magnesium carbonate is an ingredient of toothpaste.
- Calcium is used in the extraction of metals from oxides which are difficult to reduce with carbon

ANOMALOUS BEHAVIOUR OF BERYLLIUM

- Beryllium has exceptionally small atomic and ionic sizes and thus does not compare well with other members of the group
- •Because of high ionisation enthalpy and small size it forms compounds which are largely covalent and get easily hydrolysed.

ANOMALOUS BEHAVIOUR OF BERYLLIUM

- Beryllium does not exhibit coordination number more than four as in its valence shell there are only four orbitals. The remaining members of the group can have a coordination number of six by making use of d-orbitals.
- The oxide and hydroxide of beryllium, unlike the hydroxides of other elements in the group, are amphoteric in nature.

Diagonal Relationship between Beryllium and Aluminium

•The ionic radius of Be²⁺ is estimated to be 31 pm; the charge/radius ratio is nearly the same as that of the Al³⁺ ion. Hence beryllium resembles aluminium in some ways.

Similarities between

Beryllium and Aluminium

- (i) Like aluminium, beryllium is not readily attacked by acids because of the presence of an oxide film on the surface of the metal.
- (ii) Beryllium hydroxide dissolves in excess of alkali to give a beryllate ion, [Be(OH)₄] ²⁻ just as aluminium hydroxide gives aluminate ion,

- (iii) The chlorides of both beryllium and aluminium have Cl-bridged chloride structure in vapour phase. Both the chlorides are soluble in organic solvents and are strong Lewis acids. They are used as Friedel Craft catalysts.
- (iv) Beryllium and aluminium ions have strong tendency to form complexes, BeF₄²⁻, AlF₆³⁻

SOME IMPORTANT COMPOUNDS OF CALCIUM

Important compounds of calcium are

- calcium oxide,
- calcium hydroxide,
- calcium sulphate,
- calcium carbonate and cement.

These are industrially important compounds..

Calcium Oxide or Quick Lime, CaO

Preparation

• It is prepared on a commercial scale by heating limestone (CaCO₃) in a rotary kiln at 1070-1270 K.

CaCO₃
$$\xrightarrow{heat}$$
 CaO + CO₂

 The carbon dioxide is removed as soon as it is produced to enable the reaction to proceed to completion.

Properties

- Calcium oxide is a white amorphous solid.
- It has a melting point of 2870 K.
- On exposure to atmosphere, it absorbs moi

$$CaO + H_2O \rightarrow Ca(OH)_2$$

 $CaO + CO_2 \rightarrow CaCO_3$

- The addition of limited amount of water breaks the lump of lime.
- This process is called slaking of lime.

Quick lime slaked with soda gives solid **sodalime**. Being a basic oxide, it combines with acidic oxides at high temperature.

CaO + SiO₂
$$\rightarrow$$
 CaSiO₃
6CaO + P₄O₁₀ \rightarrow 2Ca₃(PO₄)₂

Uses:

• (i) It is an important primary material for manufacturing cement and is the cheapest form of alkali.

• (ii) It is used in the manufacture of sodium carbonate from caustic soda.

• (iii) It is employed in the purification of sugar and in the manufacture of dye stuffs.

Calcium Hydroxide

(Slaked lime), Ca(OH),

- Calcium hydroxide is prepared by adding water to quick lime, CaO.
- It is a white amorphous powder. It is sparingly soluble in water.
- The aqueous solution is known as lime water and a suspension of slaked lime in water is known as milk of lime.
- When carbon dioxide is passed through lime water it turns milky due to the formation of calcium carbonate.

$$Ca(OH)_2 + CO_2 \rightarrow CaCO_3 + H_2O$$

On passing excess of carbon dioxide, the precipitate dissolves to form calcium hydrogencarbonate.

$$CaCO_3 + CO_2 + H_2O \rightarrow Ca(HCO_3)_2$$

Milk of lime reacts with chlorine to form hypochlorite, a constituent of bleaching powder.

$$2Ca(OH)_2 + 2Cl_2 \rightarrow CaCl_2 + Ca(OCl)_2 + 2H_2O$$

Blanching powder

Uses:

• (i) It is used in the preparation of mortar, a building material.

• (ii) It is used in white wash due to its disinfectant nature.

• (iii) It is used in glass making, in tanning industry, for the preparation of bleaching powder and for purification

of sugar.

Calcium Carbonate, CaCO₃

- Calcium carbonate occurs in nature in several forms like limestone, chalk, marble etc.
- It can be prepared by passing carbon dioxide through slaked lime or by the addition of sodium carbonate to calcium chloride.

$$Ca(OH)_2 + CO_2 \rightarrow CaCO_3 + H_2O$$

 $CaCl_2 + Na_2CO_3 \rightarrow CaCO_3 + 2NaCl$

- Excess of carbon dioxide should be avoided since this leads to the formation of water soluble calcium hydrogencarbonate.
- Calcium carbonate is a white fluffy powder.
- It is almost insoluble in water. When heated to 1200 K, it decomposes to evolve carbon dioxide.

$$CaCO_3 \xrightarrow{1200K} CaO + CO_2$$

 It reacts with dilute acid to liberate carbon dioxide

$$CaCO_3 + 2HCl \rightarrow CaCl_2 + H_2O + CO_2$$

 $CaCO_3 + H_2SO_4 \rightarrow CaSO_4 + H_2O + CO_2$

Uses:

- It is used as a building material in the form of marble and in the manufacture of quick lime.
- Calcium carbonate along with magnesium carbonate is used as a flux in the extraction of metals such as iron.
- Specially precipitated CaCO₃ is extensively used in the manufacture of high quality paper.
- It is also used as an antacid, mild abrasive in tooth paste, a constituent of chewing

5-Mar-2gum, and a filler in cosmetics.

BIOLOGICAL IMPORTANCE OF MAGNESIUM

AND CALCIUM

- An adult body contains about 25 g of Mg and 1200 g of Ca compared with only 5 g of iron and 0.06 g of copper. The daily requirement in the human body has been estimated to be 200 300 mg.
- All enzymes that utilise ATP in phosphate transfer require magnesium as the cofactor.
- The main pigment for the absorption of light in plants is chlorophyll which contains magnesium.

- About 99 % of body calcium is present in bones and teeth.
- It also plays important roles in neuromuscular function, interneuronal transmission, cell membrane integrity and blood coagulation.
- The calcium concentration in plasma is regulated at about 100 mgL⁻¹. It is maintained by two hormones: calcitonin and parathyroid hormone.
- Bone is not an inert and unchanging substance but is continuously being solubilised and redeposited to the extent of 400 mg per day in man? All this calcium

5-Mar-2passes through the plasma.

Next chapter "P" block elements

Thank you