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# Complete CHEMISTRY

### IIT-JEE · NEET · CBSE eBOOKS CLASS 11&12th



## CLASS 11th S-Block Elements

#### 01. Introduction

S.No.	Properties	Alkaline earth metals	Alkali metals
(i)	Electronic configuration	Two electrons are present in the valency shell. The configuration is $ns^2$ .	One electron is present in the valency shell. The configuration of $ns^{1}$ .
(ii)	Valency	Bivalent	Monovalent
(iii)	Electropositive nature	Less electropositive	More electropositive.
(iv)	Hydroxides	Weak bases less soluble and decompose on heating.	Strong bases, highly soluble and stable towards heat.
(v)	Bicarbonates	These are not known in free state. Exist only in solution.	These are known in solid state.
(vi)	Carbonates	Insoluble in water. Decompose on heating.	Soluble in water. Do not decompose on heating (Li <sub>2</sub> CO <sub>3</sub> is an exception).
(vii)	Action of nitrogen	Directly combine with nitrogen and form nitrides.	Do not directly combine with nitrogen.
(viii)	Action of carbon	Directly combine with carbon and form carbides.	Do not directly combine with carbon.
(ix)	Nitrates	Decompose on heating evolving a mixture of $NO_2$ and oxygen.	Decompose on heating evolving only oxygen.
(x)	Solubility of salts	Sulphates, phosphates, fluorides, chromates, oxalates, etc. are insoluble in water.	Sulphates, phosphates, fluorides, chromates, oxalates, etc. are soluble in water.
(xi)	Physical properties	Comparatively harder. High melting points. Diamagnetic.	Soft, low melting points, Paramagnetic.
(xii)	Hydration of compounds	The compounds are extensively hydrated MgCl <sub>2</sub> . 2H <sub>2</sub> O, BaCl <sub>2</sub> .2H <sub>2</sub> O are hydrated chlorides.	The compounds are less hydrated, NaCl, KCl, RbCl form non-hydrated chlorides.
(xiii)	Reducing power	Weak as ionisation potential values are high and oxidation potential values are low.	Strong, as ionisation potential values are low and oxidation potential values are high.

Potassium, rubidium and caesium can be produced by chemical reductions.

Na(l) + KCl(l)  $\stackrel{900^{\circ}C}{=}$  NaCl(l) + K(g) Ca(s) + 2RbCl(s)  $\stackrel{\Delta}{=}$  CaCl<sub>2</sub>(s) + 2Rb(g) Mg(s) + CsOH(s)  $\stackrel{\Delta}{=}$  Mg(OH)<sub>2</sub>(s) + 2Cs(g) Ma(a) + RbOU(a)  $\stackrel{\Delta}{=}$  Ma(OH) (a) + 2Rb(a)

$$Mg(s) + RbOH(s) \rightleftharpoons Mg(OH)_2(s) + 2Rb(g)$$

These reactions occur in the direction indicated only because the metal formed escape as a gas out of the reaction mixture. Rubidium and caesium can even be obtained by reduction of their carbonates with C and Mg (in a current of  $H_2$ ) respectively.

$$RbCO_3(s) + 2C(s) \xrightarrow{\Delta} 2Rb(g) + CO(g)$$

$$CsCO_3(s) + 2C(s) \xrightarrow{\Pi_2/\Delta} MgCO_3(s) + 2Cs(g)$$

In order to get simple oxides of alkali metals like Na, K, Rb, Cs it is necessary to reduce some alkali metal compound such as nitrate. For example,

 $2\mathrm{KNO}_3(s) + 10\mathrm{K}(s) \rightarrow 6\mathrm{K}_2\mathrm{O}(s) + \mathrm{N}_2(g)$ 

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Insoluble compounds of sodium sith Si, O and Al are present in many clays. Zeolite and permutit are insoluble complex compounds containing sodium ions.

**S-Block Elements** 

Superoxides react with water to give  $\mathrm{H}_2\mathrm{O}$  and  $\mathrm{O}_2$ 

#### $2MO_2 + 2H_2O \rightarrow 2M^+ + 2OH^- + H_2O_2 + O_2$

The solubility of alkali metals in liquid ammonia may be as high as 5M. Such solutions are called expanded metals. This is because when alkali metals are dissolved in liquid ammonia, there is a considerable expansion in total volume.

When alkali metals are dissolved in liquid ammonia, a blue coloured solution is obtained. It is paramagnetic in nature. However, on increasing the concentration of alkali metal a bronze-coloured solution is obtained. This solution is diamagnetic in nature. These solutions are meta-stable and the alkali metal solwly reacts with NH<sub>3</sub> to give amide and H<sub>2</sub>.

 $2M^+$  (am) +  $2e^-$  (am) +  $2NH_3(l) \rightarrow 2MNH_2(am) + H_2(g)$ 

(M = alkali metal, am = ammonia)

#### e.g., $2Na^+$ (am) + $2e^-$ (am) + $2NH_3(l) \rightarrow 2NaNH_2(am) + H_2(g)$

However under anhydrous conditions and in the absence of catalytic impurities (such as transition metal ions), these solutions can be stored for several days.

Heavier alkali metal hydroxides form numerous hydrates e.g. NaOH,  $nH_2O$  (where n = 1, 2,

3, 4, 5 and 7). Although Li has the most negative  $E_{Li^+/Li}$  (*i.e.* highest oxidation potential) its

reaction with water is considerably less vigorous than that of sodium which has the least negative  $E^{\circ}$  among the alkali metals. Other metals react explosively with water.

All the alkali metal halides have high enthalpies of formation  $(\Delta H^{\circ}_{f})$ .

- $\Delta H_f^{\circ}$  vlaues for fluorides become less negative as we move down the group.
- $\Delta H_{f}^{\circ}$  values for fluorides, bromides and iodides become more negative as we move down the group, *e.g.*,  $\Delta H_{f}^{\circ}(\text{NaCl}) = -411.15 \text{ kJ mol}^{-1}$ ,  $\Delta H_{f}^{\circ}(\text{KCl}) = -435.9 \text{ kJ mol}^{-1}$ ,
- For a given alkali metal,  $\Delta H^{\circ}_{f}$  values always become less negative from F<sup>-</sup> to I<sup>-</sup>.  $\Delta H^{\circ}_{f}$  (kJ mol<sup>-1</sup>); NaF (-573.65); NaCl (-411.15); NaBr (-361.06); NaI (-287.78).

Melting and boiling points of halides always follow the trend

#### fluoride > chloride > bromide > iodide.

In water, the low solubility of LiF is due to its high lattice energy and the low solubility of CsI in water is due to smaller hydration energies of its two ions. Other halides of Li (C $\Gamma$ , B $r^-$  and  $\Gamma$ ) are soluble in ethyl alcohol, acetone and ethyl acetate. LiCl is soluble in pyridine. Lithium is least reactive but strongest reducing agent among all the alkali metals. Lithium unlike other alkali metals forms no acetylide on reaction with acetylene.

- Both LiCl and MgCl<sub>2</sub> are soluble in ethanol.
- Both lithium perchlorate and magnesium perchlorate are extremely soluble in alcohol.
- Both LiCl<sub>2</sub> and MgCl<sub>2</sub> are deliquescent and crystallise from aqueous solution as hydrates Licl.2H<sub>2</sub>O. MgCl<sub>2</sub>.8H<sub>2</sub>O.
- Solid bicarbonates are not formed by Li and Mg.

In the Castner-Kellner cell, during the electrolysis of aqueous NaCl, Na is preferentially discharged at the mercury cathode forming sodium amalgam. This is due to the fact that hydrogen has a high over voltage at mercury cathode. *Nafion* membrane is now widely used in the membrane cell for the manufacture of NaOH and  $Cl_2$  from NaCl(*aq*) by electrolysis. *Nafion* is a copolymer of tetrafluoromethylene and pentafluorosulphonyl ethoxy ether. The copolymer is supported by a Teflon mesh. Hydroxides of Al, Zn, Pb and Sn dissolve in excess of NaOH.