

## Solutions

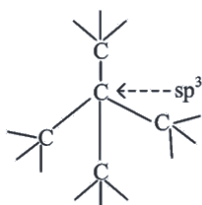
**S1.** Ans.(d)

$\text{LaH}_{2.87} \rightarrow$  Non-stoichiometric  
 $\rightarrow$  Metallic/Interstitial hydride.

**S2.** Ans.(b)

Coke: It is used as reducing agent in carbon reduction methods. (in metallurgical process)

Diamond: It is a allotrope of carbon in which each carbon is  $\text{sp}^3$  hybridised.



Fullerene: It contains pentagonal & hexagonal rings (cage like structure)

Graphite: It is soft solid because graphite layers are bonded with weak Vander Wall attractions.

**S3.** Ans.(b)

$\text{F}^{-1}$ ,  $\text{N}^{-3}$ ,  $\text{Na}^{+}$  and  $\text{O}^{-2}$

all ions are isoelectronic containing  $10e^{-}$

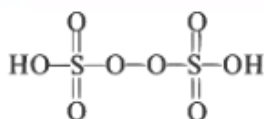
$Z_{\text{eff}} \rightarrow \text{Na}^{+} > \text{F}^{-} > \text{O}^{-2} > \text{N}^{-3}$

order of radius  $\rightarrow \text{N}^{-3} > \text{O}^{-2} > \text{F}^{-} > \text{Na}^{+}$

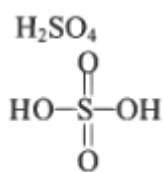
$\rightarrow$  Nitrogen to achieve Noble gas configuration by gaining  $3e^{-}$  and form  $\text{N}^{-3}$

**S4.** Ans.(a)

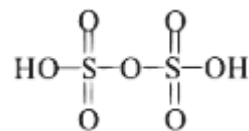
A  $\rightarrow$  Peroxodisulphuric acid



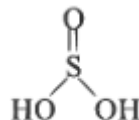
B  $\rightarrow$  Sulphuric acid



C  $\rightarrow$  Pyrosulphuric acid  $\text{H}_2\text{S}_2\text{O}_7$



D  $\rightarrow$  Sulphurous acid  $\text{H}_2\text{SO}_3$



**S5.** Ans.(b)

Borax:

$\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O} = \text{Na}_2[\text{B}_4\text{O}_5(\text{OH})_4] \cdot 8\text{H}_2\text{O}$

Kernite:  $\text{Na}_2\text{B}_4\text{O}_7 \cdot 4\text{H}_2\text{O}$

Orthoboric acid:  $\text{H}_3\text{BO}_3 = \text{B}(\text{OH})_3$

Borax bead:  $\text{NaBO}_2$  (Sodium metaborate)

**S6.** Ans.(c)

$4\text{NH}_3 + 5\text{O}_2$  (from air)  $\xrightarrow[500\text{K}, 9\text{bar}]{\text{Pt}}$   $4\text{NO} + 6\text{H}_2\text{O}$

$2\text{NO}(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2\text{NO}_2(\text{g})$

$3\text{NO}_2(\text{g}) + \text{H}_2\text{O}(\text{l}) \rightarrow 2\text{HNO}_3(\text{aq}) + \text{NO}(\text{g})$

This is industrial method of preparation of nitric acid.

**S7.** Ans.(c)

$\text{Na}_2\text{B}_4\text{O}_7 \xrightarrow{\Delta} \text{B}_2\text{O}_3 + 2\text{NaBO}_2$

Product X is  $\text{B}_2\text{O}_3$

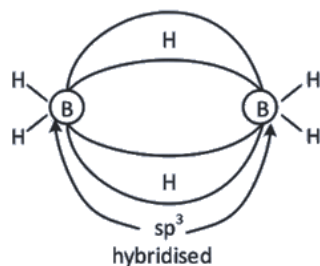
**S8.** Ans.(a)

List I (Hydrides)		List II (Nature)	
A.	$\text{MgH}_2$	$\rightarrow$	Ionic
B.	$\text{GeH}_4$	$\rightarrow$	Electron precise
C.	$\text{B}_2\text{H}_6$	$\rightarrow$	Electron deficient
D.	$\text{HF}$	$\rightarrow$	Electron rich

A-IV, B-I, C-II, D-III

**S9.** Ans.(d)

Each boron atoms in diborane uses  $\text{sp}^3$  hybrid orbitals for bonding.



**S10.** Ans.(b)

Compound	Boiling point (K)
H <sub>2</sub> O	373
H <sub>2</sub> S	213
H <sub>2</sub> Se	232
H <sub>2</sub> Te	269

The boiling points of these hybrids not exactly increases with increase in molar mass.

H<sub>2</sub>O has maximum boiling point due to intermolecular hydrogen bonding.

**S11.** Ans.(c)

Dry ice, CO<sub>2</sub>(s), is used as refrigerant

C<sub>60</sub> contains 20 six membered rings, 12 five membered rings statement 3 and 4 are correct.

**S12.** Ans.(a)

CO is Neutral oxide

BaO is Basic oxide

Al<sub>2</sub>O<sub>3</sub> is Amphoteric oxide

Cl<sub>2</sub>O<sub>7</sub> is Acidic oxide

**S13.** Ans.(b)

Carbon monoxide binds with Hb to form 300 times more stable compound carboxyhaemoglobin than oxyhaemoglobin complex.

**S14.** Ans.(d)

CO<sub>2</sub> : acidic

SnO<sub>2</sub> : amphoteric

SiO<sub>2</sub> : acidic

GeO<sub>2</sub> : acidic

**S15.** Ans.(d)

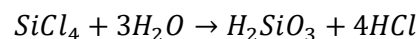
Due to presence of vacant d-orbital in si, Ge and Sn they form species like  $SiF_6^{2-}$ ,  $[GeCl_6]^{2-}$ ,  $[Sn(OH)_6]^{2-}$ .

$SiCl_6^{2-}$  does not exist because six large chloride ions cannot be accommodated around Si<sup>4+</sup> due to its small size.

**S16.** Ans.(a)

PbF<sub>4</sub> is ionic in nature, because cation is bigger and anion is smaller.

SiF<sub>4</sub> is easily hydrolysed because it has unoccupied 3d orbital that are able to accept electron pairs from the oxygen atoms on water to form bond.



GeX<sub>4</sub> is more stable than GeX<sub>2</sub> as in GeX<sub>4</sub> all the orbitals are fully filled.

SnF<sub>4</sub> is ionic in nature as F atom is very small and Sn atom is very large. So, it is ionic in nature according to Fajan's rule.

**S17.** Ans.(c)

Group 13 order of atomic radius is not regular due to transition contraction.

So order is: B < Ga < Al < In < Tl

**S18.** Ans.(d)

∴ 'B' has no vacant d-orbitals in its valence shell, so it can't extend its covalency beyond 4, i.e., 'B' cannot form the ion like  $MF_6^{3-}$  i.e.  $BF_6^{3-}$ .

Hence, the correct option is (d).

**S19.** Ans.(b)

Sn<sup>2+</sup> is reducing while Pb<sup>4+</sup> is oxidizing. This is because of absence of f-orbital (fully filled) in Sn<sup>2+</sup> and presence in Pb<sup>4+</sup> due to which Pb<sup>4+</sup> show inert pair effect but not Sn<sup>2+</sup>.

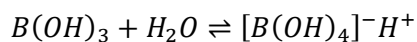
**S20.** Ans.(b)

+1 oxidation state increases down the group as the fully filled d- and f- orbitals will get added due to which inert pair effect will come into role.

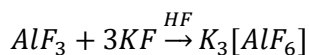
Tl > In > Ga > Al > B

**S21.** Ans.(a)

Boric acid is a Lewis acid, it accepts a pair of electron in aqueous solution to complete its octet because it is an electron deficient compound.



**S22.** Ans.(b)



$AlF_3$  is insoluble in Anhydrous HF because  $F^-$  ion are not available in Hydrogen bonded HF.

**S23.** Ans.(c)

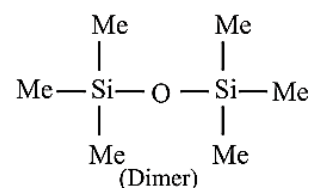
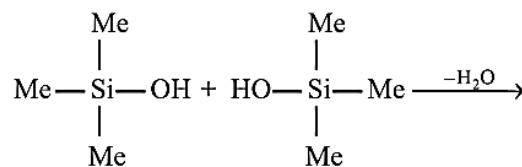
Order of oxidation state in group 13 is  $Al < Ga < In < Tl$

(+1) oxidation state stability increases down the group because of inert pair effect.

**S24.** Ans.(c)

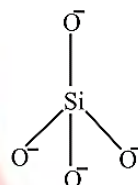
$Me_3SiCl$  is not a monomer for high molecular mass silicone polymer because it generates  $Me_3SiOH$  when subjected to hydrolysis which contains only one reacting site.

Hence, the polymerization stops just after first step.



**S25.** Ans.(b)

Basic unit for silicates is  $SiO_4^{4-}$



**S26.** Ans.(a)

BN is also known as inorganic graphite with  $sp^2$  hybridisation and hexagonal structure.

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