# **NCERT Solutions for Class 11 Chemistry Chapter 11**

# The p-Block Elements Class 11

Chapter 11 The p-Block Elements Exercise Solutions

Exercise: Solutions of Questions on Page Number: 323

Q1:

Discuss the pattern of variation in the oxidation states of

(i) B to TI and (ii) C to Pb.

Answer:

(i) B to TI

The electric configuration of group 13 elements is  $ns^2$   $np^1$ . Therefore, the most common oxidation state exhibited by them should be +3. However, it is only boron and aluminium which practically show the +3 oxidation state. The remaining elements, i.e., Ga, In, TI, show both the +1 and +3 oxidation states. On moving down the group, the +1 state becomes more stable. For example, TI (+1) is more stable than TI (+3). This is because of the inert pair effect. The two electrons present in the s-shell are strongly attracted by the nucleus and do not participate in bonding. This inert pair effect becomes more and more prominent on moving down the group. Hence, Ga (+1) is unstable, In (+1) is fairly stable, and TI (+1) is very stable.

Group 13 element	Oxidation state
В	+3
Al	+3
Ga, In, Tl	+1, +3

The stability of the +3 oxidation state decreases on moving down the group.

# (ii) C to Pb

The electronic configuration of group 14 elements is  $ns^2 np^2$ . Therefore, the most common oxidation state exhibited by them should be +4. However, the +2 oxidation state becomes more and more common on moving down the group. C and Si mostly show the +4 state. On moving down the group, the higher oxidation state becomes less stable. This is because of the inert pair effect. Thus, although Ge, Sn, and Pb show both the +2 and +4 states, the stability of the lower oxidation state increases and that of the higher oxidation state decreases on moving down the group.

Group 14 element	Oxidation state
С	+4
Si	+4
Ge, Sn, Pb	+2, +4

# Q2:

How can you explain higher stability of BCI<sub>3</sub> as compared to TICI<sub>3</sub>?

### Answer:

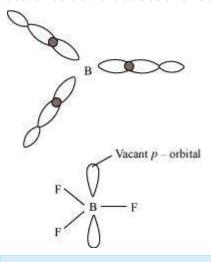
Boron and thallium belong to group 13 of the periodic table. In this group, the +1 oxidation state becomes more stable on moving down the group. BCI<sub>3</sub>is more stable than TICI<sub>3</sub>because the +3 oxidation state of B is more stable than the +3 oxidation state of TI. In TI, the +3 state is highly oxidising and it reverts back to the more stable +1 state.

# Q3:

Why does boron trifluoride behave as a Lewis acid?

### Answer:

The electric configuration of boron is  $ns^2 np^1$ . It has three electrons in its valence shell. Thus, it can form only three covalent bonds. This means that there are only six electrons around boron and its octet remains incomplete. When one atom of boron combines with three fluorine atoms, its octet remains incomplete. Hence, boron trifluoride remains electron-deficient and acts as a Lewis acid.



# Q4:

Consider the compounds, BCI<sub>3</sub> and CCI<sub>4</sub>. How will they behave with water? Justify.

### Answer:

Being a Lewis acid, BCl<sub>3</sub> readily undergoes hydrolysis. Boric acid is formed as a result.

$$BCl_3 + 3H_2O \longrightarrow 3HCl + B(OH)_3$$

CCl<sub>4</sub>completely resists hydrolysis. Carbon does not have any vacant orbital. Hence, it cannot accept electrons from water to form an intermediate. When CCl<sub>4</sub>and water are mixed, they form separate layers.

### Q5:

Is boric acid a protic acid? Explain.

#### Answer:

Boric acid is not a protic acid. It is a weak monobasic acid, behaving as a Lewis acid.

$$B(OH)_3 + 2HOH \longrightarrow [B(OH)_4]^- + H_3O^+$$

It behaves as an acid by accepting a pair of electrons from a OH ion.

# Q6:

Explain what happens when boric acid is heated.

# Answer:

On heating orthoboric acid ( $H_3BO_3$ ) at 370 K or above, it changes to metaboric acid ( $HBO_2$ ). On further heating, this yields boric oxide  $B_2O_3$ .

$$H_3BO_3 \xrightarrow{\Delta \atop 370 \text{ K}} HBO_2 \xrightarrow{\Delta \atop \text{red hot}} B_2O_3$$
Metaboric acid Boric oxide

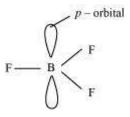
### Q7:

Describe the shapes of BF<sub>3</sub> and BH<sub>4</sub>. Assign the hybridisation of boron in these species.

# Answer:

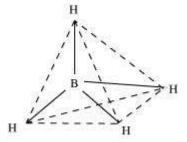
### (i) BF<sub>3</sub>

As a result of its small size and high electronegativity, boron tends to form monomeric covalent halides. These halides have a planar triangular geometry. This triangular shape is formed by the overlap of three  $sp^2$ hybridised orbitals of boron with the sporbitals of three halogen atoms. Boron is  $sp^2$ hybridised in BF<sub>3</sub>.



# (ii) BH<sub>4</sub>

Boron-hydride ion (BH<sub>4</sub>) is formed by the  $sp^3$ hybridisation of boron orbitals. Therefore, it is tetrahedral in structure.



#### Q8:

Write reactions to justify amphoteric nature of aluminium.

### Answer:

A substance is called amphoteric if it displays characteristics of both acids and bases. Aluminium dissolves in both acids and bases, showing amphoteric behaviour.

(i) 
$$2Al_{(s)} + 6HCl_{(aq)} \longrightarrow 2Al_{(aq)}^{3+} + 6Cl_{(aq)}^{-} + 3H_{2(g)}$$

(ii) 
$$2Al_{(s)} + 2NaOH_{(aq)} + 6H_2O_{(l)} \longrightarrow 2Na^+ \left[Al(OH)_4\right]_{(aq)}^- + 3H_{2(g)}$$

### Q9:

What are electron deficient compounds? Are BCI<sub>3</sub> and SiCI<sub>4</sub> electron deficient species? Explain.

### Answer:

In an electron-deficient compound, the octet of electrons is not complete, i.e., the central metal atom has an incomplete octet. Therefore, it needs electrons to complete its octet.

# (i) BCI<sub>3</sub>

BCI<sub>3</sub>is an appropriate example of an electron-deficient compound. B has 3 valence electrons. After forming three covalent bonds with chlorine, the number of electrons around it increases to 6. However, it is still short of two electrons to complete its octet.

# (ii) SiCI<sub>4</sub>

The electronic configuration of silicon is  $ns^2 np^2$ . This indicates that it has four valence electrons. After it forms four covalent bonds with four chlorine atoms, its electron count increases to eight. Thus, SiCl<sub>4</sub>is not an electron-deficient compound.

# Q10:

Write the resonance structures of  ${}^{CO_3^{2-}}$  and  ${}^{HCO_3^{-}}.$ 

# Answer:

There are only two resonating structures for the bicarbonate ion.

# Q11:

What is the state of hybridisation of carbon in (a)  ${
m CO}_3^{2-}$  (b) diamond (c) graphite?

# Answer:

The state of hybridisation of carbon in:

(a)  $CO_3^{2-}$ 

C in  ${
m CO}_3^{2-}$  is  ${\it sp}^2$ hybridised and is bonded to three oxygen atoms.

# (b) Diamond

Each carbon in diamond is *sp*<sup>3</sup> hybridised and is bound to four other carbon atoms.

# (c) Graphite

Each carbon atom in graphite is  $sp^2$ hybridised and is bound to three other carbon atoms.

# Q12:

Explain the difference in properties of diamond and graphite on the basis of their structures.

### Answer:

Diamond	Graphite
It has a crystalline lattice.	It has a layered structure.
In diamond, each carbon atom is $sp^s$ hybridised and is bonded to four other carbon atoms through a $\tilde{A}\mathcal{E}$ ' bond.	In graphite, each carbon atom is $sp^2$ hybridised and is bonded to three other carbon atoms through a $\tilde{A}\mathcal{E}$ ' bond. The fourth electron forms a $\pi$ bond.
It is made up of tetrahedral units.	It has a planar geometry.
The C–C bond length in diamond is 154 pm.	The C–C bond length in graphite is 141.5 pm.
It has a rigid covalent bond network which is difficult to break.	It is quite soft and its layers can be separated easily.
It acts as an electrical insulator.	It is a good conductor of electricity.

# Q13:

Rationalise the given statements and give chemical reactions:

"¢ Lead(II) chloride reacts with Cl₂to give PbCl₄.

"¢ Lead(IV) chloride is highly unstable towards heat.

"¢ Lead is known not to form an iodide, Pbl4.

### Answer:

(a) Lead belongs to group 14 of the periodic table. The two oxidation states displayed by this group is +2 and +4. On moving down the group, the +2 oxidation state becomes more stable and the +4 oxidation state becomes less stable. This is because of the inert pair effect. Hence, PbCl<sub>4</sub>is much less stable than PbCl<sub>2</sub>. However, the formation of PbCl<sub>4</sub>takes place when chlorine gas is bubbled through a saturated solution of PlCl<sub>2</sub>.

$$PbCl_{2(s)} + Cl_{2(g)} \longrightarrow PbCl_{4(l)}$$

**(b)** On moving down group IV, the higher oxidation state becomes unstable because of the inert pair effect. Pb(IV) is highly unstable and when heated, it reduces to Pb(II).

$$PbCl_{4(I)} \xrightarrow{\Delta} PbCl_{2(s)} + Cl_{2(g)}$$

(c) Lead is known not to form PbI<sub>4</sub>. Pb (+4) is oxidising in nature and I is reducing in nature. A combination of Pb(IV) and iodide ion is not stable. Iodide ion is strongly reducing in nature. Pb(IV) oxidises I<sup>ac</sup> to I<sup>2</sup> and itself gets reduced to Pb(II).

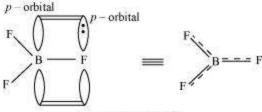
$$PbI_4 \longrightarrow PbI_7 + I_7$$

# Q14:

Suggest reasons why the B–F bond lengths in BF₃(130 pm) and BF⁴ (143 pm) differ.

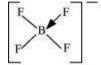
# Answer:

The Bâ $\in$ "F bond length in BF $_3$ is shorter than the Bâ $\in$ "F bond length in With a vacant p-orbital on boron, the fluorine and boron atoms undergo  $p\pi$ â $\in$ " $p\pi$ back-bonding to remove this deficiency. This imparts a double-bond character to the Bâ $\in$ "F bond.



 $p\pi$ - $p\pi$  back bonding

This double-bond character causes the bond length to shorten in BF $_3$ (130 pm). However, when BF $_3$ coordinates with the fluoride ion, a change in hybridisation from  $sp^2$ (in BF $_3$ ) to  $sp^3$ (in  $^{BF}_4^-$ ) occurs. Boron now forms  $4\tilde{A}$ /E'bonds and the double-bond character is lost. This accounts for a Bâ $\in$ "F bond length of 143 pm in  $^{BF}_4^-$  ion.



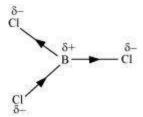
Tetrahedral structure

# Q15:

If B-CI bond has a dipole moment, explain why BCI₃molecule has zero dipole moment.

### Answer:

As a result of the difference in the electronegativities of B and CI, the B-CI bond is polar in nature. However, the BCI<sub>3</sub>molecule is non-polar. This is because BCI<sub>3</sub>is trigonal planar in shape. It is a symmetrical molecule. Hence, the respective dipole-moments of the B-CI bond cancel each other, thereby causing a zero-dipole moment.



# Q16:

Aluminium trifluoride is insoluble in anhydrous HF but dissolves on addition of NaF. Aluminium trifluoride precipitates out of the resulting solution when gaseous BF<sub>3</sub> is bubbled through. Give reasons.

### Answer:

Hydrogen fluoride (HF) is a covalent compound and has a very strong intermolecular hydrogen-bonding. Thus, it does not provide ions and aluminium fluoride (AIF) does not dissolve in it. Sodium fluoride (NaF) is an ionic compound and when it is added to the mixture, AIF dissolves. This is because of the availability of free F<sup>ac</sup>. The reaction involved in the process is:

$$AlF_3 + 3 NaF \longrightarrow Na_3[AlF_6]$$
  
Sodium hexafluoroaluminate (III)

When boron trifluoride (BF<sub>3</sub>) is added to the solution, aluminium fluoride precipitates out of the solution. This happens because the tendency of boron to form complexes is much more than that of aluminium. Therefore, when BF<sub>3</sub>is added to the solution, B replaces Al from the complexes according to the following reaction:

$$Na_3[AlF_6] + 3 BF_3 \longrightarrow 3 Na[BF_4] + AlF_3$$

### Q17:

Suggest a reason as to why CO is poisonous.

# Answer:

Carbon monoxide is highly-poisonous because of its ability to form a complex with haemoglobin. The CO-Hb complex is more stable than the  $O_2$ -Hb complex. The former prevents Hb from binding with oxygen. Thus, a person dies because of suffocation on not receiving oxygen. It is found that the

CO-Hb complex is about 300 times more stable than the O<sub>2</sub>-Hb complex.

### Q18:

How is excessive content of CO<sub>2</sub> responsible for global warming?

### Answer:

Carbon dioxide is a very essential gas for our survival. However, an increased content of CO<sub>2</sub>in the atmosphere poses a serious threat. An increment in the combustion of fossil fuels, decomposition of limestone, and a decrease in the number of trees has led to greater levels of carbon dioxide. Carbon dioxide has the property of trapping the heat provided by sunrays. Higher the level of carbon dioxide, higher is the amount of heat trapped. This results in an increase in the atmospheric temperature, thereby causing global warming.

#### Q19:

Explain structures of diborane and boric acid.

### Answer:

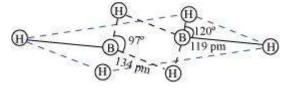
### (a) Diborane

 $B_2H_6$  is an electron-deficient compound.  $B_2H_6$  has only 12 electrons - 6 efrom 6 H atoms and 3 ereach from 2 B atoms. Thus, after combining with 3 H atoms, none of the boron atoms has any electrons left. X-ray diffraction studies have shown the structure of diborane as:

$$H_t$$
 = terminal hydrogen  
 $H_b$  = bridging hydrogen  
 $H_t$ 

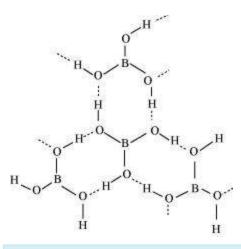
$$\begin{array}{c}
H_t \\
H_t
\end{array}$$
 $\begin{array}{c}
H_b \\
H_t
\end{array}$ 
 $\begin{array}{c}
H_t \\
H_t
\end{array}$ 

2 boron and 4 terminal hydrogen atoms ( $H_t$ ) lie in one plane, while the other two bridging hydrogen atoms ( $H_t$ ) lie in a plane perpendicular to the plane of boron atoms. Again, of the two bridging hydrogen atoms, one H atom lies above the plane and the other lies below the plane. The terminal bonds are regular two-centre two-electron ( $2c - 2e^{-}$ ) bonds, while the two bridging (B-H-B) bonds are three-centre two-electron ( $3c - 2e^{-}$ ) bonds.



# (b) Boric acid

Boric acid has a layered structure. Each planar  $BO_3$  unit is linked to one another through H atoms. The H atoms form a covalent bond with a  $BO_3$  unit, while a hydrogen bond is formed with another  $BO_3$  unit. In the given figure, the dotted lines represent hydrogen bonds.



# Q20:

What happens when

- (a) Borax is heated strongly,
- (b) Boric acid is added to water,
- (c) Aluminium is treated with dilute NaOH,
- (d) BF<sub>3</sub>is reacted with ammonia?

# Answer:

(a) When heated, borax undergoes various transitions. It first loses water molecules and swells. Then, it turns into a transparent liquid, solidifying to form a glass-like material called borax bead.

$$Na_2B_4O_7 \cdot 10H_2O \xrightarrow{\Delta} Na_2B_4O_7 \xrightarrow{\Delta} 2NaBO_2 + B_2O_3$$
  
Borax Sodium Boric metaborate anhydride

**(b)** When boric acid is added to water, it accepts electrons from <sup>a∈</sup>OH ion.

$$B(OH)_3 + 2HOH \longrightarrow [B(OH)_4]^- + H_3O^+$$

(c) Al reacts with dilute NaOH to form sodium tetrahydroxoaluminate(III). Hydrogen gas is liberated in the process.

$$2 \text{ Al}_{(s)} + 2 \text{ NaOH}_{(aq)} + 6 \text{ H}_2 \text{O}_{(l)} \longrightarrow 2 \text{ Na}^+ \left[ \text{Al}(\text{OH})_4 \right]_{(aq)}^- + 3 \text{ H}_{2(g)}$$

(d) BF<sub>3</sub>(a Lewis acid) reacts with NH<sub>3</sub>(a Lewis base) to form an adduct. This results in a complete octet around B in BF<sub>2</sub>

$$F_3B + :NH_3 \longrightarrow F_3B \leftarrow :NH_3$$

### **Explain the following reactions**

- (a) Silicon is heated with methyl chloride at high temperature in the presence of copper;
- (b) Silicon dioxide is treated with hydrogen fluoride;
- (c) CO is heated with ZnO;
- (d) Hydrated alumina is treated with aqueous NaOH solution.

#### Answer:

(a) When silicon reacts with methyl chloride in the presence of copper (catalyst) and at a temperature of about 537 K, a class of organosilicon polymers called methyl-substituted chlorosilanes (MeSiCl<sub>3</sub>, Me<sub>2</sub>SiCl<sub>2</sub>, Me<sub>3</sub>SiCl, and Me<sub>4</sub>Si) are formed.

**(b)** When silicon dioxide (SiO₂) is heated with hydrogen fluoride (HF), it forms silicon tetrafluoride (SiF₄). Usually, the Si–O bond is a strong bond and it resists any attack by halogens and most acids, even at a high temperature. However, it is attacked by HF.

$$SiO_2 + 4HF \longrightarrow SiF_4 + 2H_2O$$

The SiF₄formed in this reaction can further react with HF to form hydrofluorosilicic acid.

$$SiF_4 + 2HF \longrightarrow H_2SiF_6$$

(c) When CO reacts with ZnO, it reduces ZnO to Zn. CO acts as a reducing agent.

$$ZnO_{(s)} + CO_{(g)} \xrightarrow{\Delta} Zn_{(s)} + CO_{2(g)}$$

(d) When hydrated alumina is added to sodium hydroxide, the former dissolves in the latter because of the formation of sodium meta-aluminate.

$$Al_2O_3 \cdot 2H_2O + 2 \text{ NaOH} \longrightarrow 2 \text{ NaAlO}_3 + 3H_2O$$

# Q22:

### Give reasons:

- (i) Conc. HNO<sub>3</sub>can be transported in aluminium container.
- (ii) A mixture of dilute NaOH and aluminium pieces is used to open drain.
- (iii) Graphite is used as lubricant.
- (iv) Diamond is used as an abrasive.
- (v) Aluminium alloys are used to make aircraft body.
- (vi) Aluminium utensils should not be kept in water overnight.
- (vii) Aluminium wire is used to make transmission cables.

### Answer:

- (i) Concentrated HNO₃can be stored and transported in aluminium containers as it reacts with aluminium to form a thin protective oxide layer on the aluminium surface. This oxide layer renders aluminium passive.
- (ii) Sodium hydroxide and aluminium react to form sodium tetrahydroxoaluminate(III) and hydrogen gas. The pressure of the produced hydrogen gas is used to open blocked drains.

$$2 \text{ Al} + 2 \text{ NaOH} + 6 \text{ H,O} \longrightarrow 2 \text{ Na}^{+} [\text{Al(OH)}_{4}]^{+} + 3 \text{ H},$$

- (iii) Graphite has a layered structure and different layers of graphite are bonded to each other by weak van der Waals' forces. These layers can slide over each other. Graphite is soft and slippery. Therefore, graphite can be used as a lubricant.
- (iv) In diamond, carbon is  $sp^3$ hybridised. Each carbon atom is bonded to four other carbon atoms with the help of strong covalent bonds. These covalent bonds are present throughout the surface, giving it a very rigid 3-D structure. It is very difficult to break this extended covalent bonding and for this reason, diamond is the hardest substance known. Thus, it is used as an abrasive and for cutting tools.
- (v) Aluminium has a high tensile strength and is very light in weight. It can also be alloyed with various metals such as Cu, Mn, Mq, Si, and Zn. It is very malleable and ductile. Therefore, it is used in making aircraft bodies.
- (vi) The oxygen present in water reacts with aluminium to form a thin layer of aluminium oxide. This layer prevents aluminium from further reaction. However, when water is kept in an aluminium vessel for long periods of time, some amount of aluminium oxide may dissolve in water. As aluminium ions are harmful, water should not be stored in aluminium vessels overnight.
- (vii) Silver, copper, and aluminium are among the best conductors of electricity. Silver is an expensive metal and silver wires are very expensive. Copper is quite expensive and is also very heavy. Aluminium is a very ductile metal. Thus, aluminium is used in making wires for electrical conduction.

### Q23:

Explain why is there a phenomenal decrease in ionisation enthalpy from carbon to silicon?

Answer:

Ionisation enthalpy of carbon (the first element of group 14) is very high (1086 kJ/mol). This is expected owing to its small size. However, on moving down the group to silicon, there is a sharp decrease in the enthalpy (786 kJ). This is because of an appreciable increase in the atomic sizes of elements on moving down the group.

### Q24:

How would you explain the lower atomic radius of Ga as compared to AI?

# Answer:

Atomic radius (in pm)	
Aluminium	143
Gallium	135

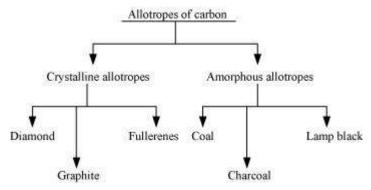
Although Ga has one shell more than Al, its size is lesser than Al. This is because of the poor shielding effect of the 3*d*-electrons. The shielding effect of *d*-electrons is very poor and the effective nuclear charge experienced by the valence electrons in gallium is much more than it is in the case of Al.

# Q25:

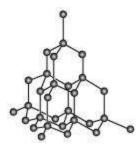
What are allotropes? Sketch the structure of two allotropes of carbon namely diamond and graphite. What is the impact of structure on physical properties of two allotropes?

### Answer:

Allotropy is the existence of an element in more than one form, having the same chemical properties but different physical properties. The various forms of an element are called allotropes.

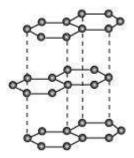


# Diamond:



The rigid 3-D structure of diamond makes it a very hard substance. In fact, diamond is one of the hardest naturally-occurring substances. It is used as an abrasive and for cutting tools.

# **Graphite:**



It has *sp*<sup>2</sup>hybridised carbon, arranged in the form of layers. These layers are held together by weak van der Walls' forces. These layers can slide over each other, making graphite soft and slippery. Therefore, it is used as a lubricant.

# Q26:

(a) Classify following oxides as neutral, acidic, basic or amphoteric:

CO, B<sub>2</sub>O<sub>3</sub>, SiO<sub>2</sub>, CO<sub>2</sub>, Al<sub>2</sub>O<sub>3</sub>, PbO<sub>2</sub>, Tl<sub>2</sub>O<sub>3</sub>

(b) Write suitable chemical equations to show their nature.

# Answer:

- (1) CO = Neutral
- (2)  $B_2O_3$ = Acidic

Being acidic, it reacts with bases to form salts. It reacts with NaOH to form sodium metaborate.

$$B_2O_3 + 2 NaOH \longrightarrow 2 NaBO_2 + H_2O$$

(3)  $SiO_2 = Acidic$ 

Being acidic, it reacts with bases to form salts. It reacts with NaOH to form sodium silicate.

$$SiO_{2} + 2 NaOH \longrightarrow 2 Na_{2}SiO_{3} + H_{2}O$$

(4) CO<sub>2</sub>= Acidic

Being acidic, it reacts with bases to form salts. It reacts with NaOH to form sodium carbonate.

$$CO_2 + 2 NaOH \longrightarrow Na_2CO_3 + H_2O$$

(5) Al<sub>2</sub>O<sub>3</sub>= Amphoteric

Amphoteric substances react with both acids and bases. Al<sub>2</sub>O<sub>3</sub>reacts with both NaOH and H<sub>2</sub>SO<sub>4</sub>.

$$Al_2O_3 + 2 NaOH \longrightarrow NaAlO_3$$

$$Al_2O_3 + 3H_2SO_4 \longrightarrow Al_2(SO_4)_3 + 3H_2O$$

(6) PbO<sub>2</sub>= Amphoteric

Amphoteric substances react with both acids and bases. PbO2reacts with both NaOH and H2SO4.

$$PbO_2 + 2 NaOH \longrightarrow Na_2 PbO_3 + H_2O$$

$$2 \text{ PbO}_2 + 2 \text{ H}_2 \text{SO}_4 \longrightarrow 2 \text{ PbSO}_4 + 2 \text{ H}_2 \text{O} + \text{O}_2$$

(7) Tl<sub>2</sub>O<sub>3</sub>= Basic

Being basic, it reacts with acids to form salts. It reacts with HCl to form thallium chloride.

$$Tl_2O_3 + 6HCl \longrightarrow 2TlCl_3 + 3H_2O$$

### Q27:

In some of the reactions thallium resembles aluminium, whereas in others it resembles with group I metals. Support this statement by giving some evidences.

### Answer:

Thallium belongs to group 13 of the periodic table. The most common oxidation state for this group is +3. However, heavier members of this group also display the +1 oxidation state. This happens because of the inert pair effect. Aluminium displays the +3 oxidation state and alkali metals display the +1 oxidation state. Thallium displays both the oxidation states. Therefore, it resembles both aluminium and alkali metals.

Thallium, like aluminium, forms compounds such as TICl<sub>3</sub>and Tl<sub>2</sub>O<sub>3</sub>. It resembles alkali metals in compounds Tl<sub>2</sub>O and TICl.

### Q28:

When metal X is treated with sodium hydroxide, a white precipitate (A) is obtained, which is soluble in excess of NaOH to give soluble complex (B). Compound (A) is soluble in dilute HCl to form compound (C). The compound (A) when heated strongly gives (D), which is used to extract metal. Identify (X), (A), (B), (C) and (D). Write suitable equations to support their identities.

# Answer:

The given metal X gives a white precipitate with sodium hydroxide and the precipitate dissolves in excess of sodium hydroxide. Hence, X must be aluminium.

The white precipitate (compound A) obtained is aluminium hydroxide. The compound B formed when an excess of the base is added is sodium tetrahydroxoaluminate(III).

2Al + 3 NaOH 
$$\longrightarrow$$
 A  $I(OH)_3 \downarrow + 3 Na^+$   
Aluminium (X) Sodium hydroxide White ppt.(A)

$$Al(OH)_3 + NaOH \longrightarrow Na^+[Al(OH)_4]^-$$
(A) Sodium tetrahydroxoaluminate (III)
(Soluble complex B)

Now, when dilute hydrochloric acid is added to aluminium hydroxide, aluminium chloride (compound C) is obtained.

$$Al(OH)_3 + 3 HCl \longrightarrow AlCl_3 + 3 H_2O$$
(A) (C)

Also, when compound A is heated strongly, it gives compound D. This compound is used to extract metal X. Aluminium metal is extracted from alumina. Hence, compound D must be alumina.

$$2 \text{ Al(OH)}_3 \xrightarrow{\Delta} \text{Al}_2\text{O}_3 + 3 \text{ H}_2\text{O}$$
(A) (D)

# Q29:

What do you understand by (a) inert pair effect (b) allotropy and(c) catenation?

### Answer:

### (a) Inert pair effect

As one moves down the group, the tendency of s-block electrons to participate in chemical bonding decreases. This effect is known as inert pair effect. In case of group 13 elements, the electronic configuration is  $ns^2$   $np^1$  and their group valency is +3. However, on moving down the group, the +1 oxidation state becomes more stable. This happens because of the poor shielding of the  $ns^2$  electrons by the d- and f- electrons. As a result of the poor shielding, the  $ns^2$  electrons are held tightly by the nucleus and so, they cannot participate in chemical bonding.

# (b) Allotropy

Allotropy is the existence of an element in more than one form, having the same chemical properties but different physical properties. The various forms of an element are called allotropes. For example, carbon exists in three allotropic forms: diamond, graphite, and fullerenes.

# (c) Catenation

The atoms of some elements (such as carbon) can link with one another through strong covalent bonds to form long chains or branches. This property is known as catenation. It is most common in carbon and quite significant in Si and S.

### Q30:

A certain salt X, gives the following results.

- (i) Its aqueous solution is alkaline to litmus.
- (ii) It swells up to a glassy material Yon strong heating.
- (iii) When conc. H₂SO₄is added to a hot solution of X, white crystal of an acid Zseparates out.

Write equations for all the above reactions and identify X, Y and Z.

### Answer:

The given salt is alkaline to litmus. Therefore, Xis a salt of a strong base and a weak acid. Also, when Xis strongly heated, it swells to form substance Y. Therefore, Xmust be borax.

When borax is heated, it loses water and swells to form sodium metaborate. When heating is continued, it solidifies to form a glassy material Y. Hence, Ymust be a mixture of sodium metaborate and boric anhydride.

$$Na_2B_4O_7 + 7H_2O \xrightarrow{water} 2NaOH + 4H_3BO_3$$
  
Borax (X) Sodium hydroxide Orthoboric acid

$$Na_2B_4O_7 \cdot 10H_2O \xrightarrow{\Delta} Na_2B_4O_7 \xrightarrow{\Delta} B_2O_3 + 2NaBO_2$$
  
 $Borax(X)$  Sodium metaborate Boric anhydride (Glassy material)

When concentrated acid is added to borax, white crystals of orthoboric acid (Z) are formed.

$$Na_2B_4O_7 \cdot 10H_2O + H_2SO_4 \xrightarrow{\Delta} Na_2SO_4 + 4H_3BO_3 + 5H_2O$$
  
Borax (X) Orthoboric acid (Z)

# Q31:

Write balanced equations for:

- (i) BF₃ + LiH â†'
- (ii) B₂H<sub>6</sub> + H₂O →
- (iii) NaH + B₂H<sub>6</sub> â†'
- (iv)  $H_3BO_3 \xrightarrow{\Delta}$
- (v) AI + NaOH â†'
- (vi) B<sub>2</sub>H<sub>6</sub> + NH<sub>3</sub> â†'

# Answer:

(i) 
$$2BF_3$$
 +  $6LiH$   $\longrightarrow$   $B_2H_6$  +  $6LiF$   
Boron trifluoride Lithium hydride Diborane Lithium fluoride

(ii) 
$$B_2H_6 + 6H_2O \longrightarrow 2H_3BO_3 + 6H_2$$
  
Diborane Water Orthoboric acid Hydrogen

(iii) 
$$B_2H_6 + 2NaH \xrightarrow{\text{ether}} 2NaBH_4$$
  
Diborane Sodium hydride Sodium borohydride

(iv)

$$4H_3BO_3$$
  $-4H_2O$   $-4H_2O$   $-4H_2O$   $-4H_2O$   $-4H_2O$   $-4H_2O$   $-410$  K  $-410$  K  $-410$  K  $-410$  Red hot  $-4$ 

(v) 
$$2AI + 2NaOH + 6H_2O \longrightarrow 2Na^+[AI(OH)_4]_{(aq)}^- + 3H_2$$

Sodium tetrahydroxoaluminate(III)

(vi) 
$$3 B_2 H_6 + 6 NH_3 \longrightarrow 3[BH_2(NH_3)_2]^+[BH_4]^- \longrightarrow 2B_3 N_3 H_6 + 12H_2$$
  
Borazene

# Q32:

Give one method for industrial preparation and one for laboratory preparation of CO and CO2 each.

# Answer:

# Caron dioxide

In the laboratory, CO<sub>2</sub>can be prepared by the action of dilute hydrochloric acid on calcium carbonate. The reaction involved is as follows:

$$CaCO_3 + 2HCl_{(aq)} \longrightarrow CaCl_{2(aq)} + CO_{2(g)} + H_2O_{(I)}$$

CO<sub>2</sub> is commercially prepared by heating limestone. The reaction involved is as follows:

# Caron monoxide

In the laboratory, CO is prepared by the dehydration of formic acid with conc. H<sub>2</sub>SO<sub>4</sub>, at 373 K. The reaction involved is as follows:

$$HCOOH \xrightarrow{373 \text{ K}} H_2O + CO \uparrow$$

CO is commercially prepared by passing steam over hot coke. The reaction involved is as follows:

$$C_{(s)} + H_2O_{(g)} \xrightarrow{473 - 1273 \text{ K}} \underbrace{CO_{(g)} + H_{2(g)}}_{\text{water gas}}$$

# Q33:

An aqueous solution of borax is

- (a) neutral (b) amphoteric
- (c) basic (d) acidic

# Answer:

(c) Borax is a salt of a strong base (NaOH) and a weak acid (H<sub>3</sub>BO<sub>3</sub>). It is, therefore, basic in nature.

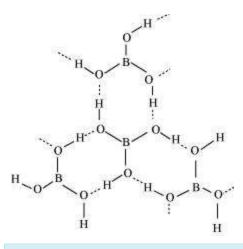
# Q34:

Boric acid is polymeric due to

- (a) its acidic nature (b) the presence of hydrogen bonds
- (c) its monobasic nature (d) its geometry

# Answer:

**(b)** Boric acid is polymeric because of the presence of hydrogen bonds. In the given figure, the dotted lines represent hydrogen bonds.



# Q35:

The type of hybridisation of boron in diborane is

(a) sp (b)  $sp^{2}$  (c)  $sp^{3}$  (d)  $dsp^{2}$ 

# Answer:

(c) Boron in diborane is  $sp^3$ hybridised.

# Q36:

Thermodynamically the most stable form of carbon is

- (a) diamond (b) graphite
- (c) fullerenes (d) coal

# Answer:

**(b)** Graphite is thermodynamically the most stable form of carbon.

# Q37:

Elements of group 14

- (a) exhibit oxidation state of +4 only
- (b) exhibit oxidation state of +2 and +4
- (c) form M<sup>2</sup>-and M⁴+ion
- (d) form M2+and M4+ions

# Answer:

**(b)**The elements of group 14 have 4 valence electrons. Therefore, the oxidation state of the group is +4. However, as a result of the inert pair effect, the lower oxidation state becomes more and more stable and the higher oxidation state becomes less stable. Therefore, this group exhibits +4 and +2 oxidation states.

Group 14 element	Oxidation state
С	+4
Si	+4
Ge, Sn, Pb	+2, +4

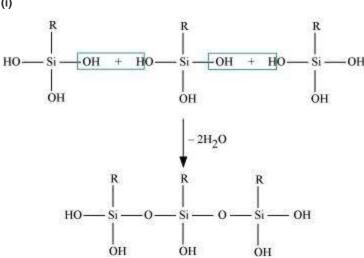
# Q38:

If the starting material for the manufacture of silicones is RSiCl<sub>3</sub>, write the structure of the product formed.

# Answer:

$$RSiCl_3 + 3H_2O \longrightarrow RSi(OH)_3 + 3HCl$$

(i)



(ii)

