

CHAPTER - 11

P-BLOCK ELEMENTS

Question 11.1:

Discuss the pattern of variation in the oxidation states of

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

Answer 11.1:

(i) B to Tl

The electronic 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, Tl, show both the +1 and +3 oxidation states. On moving down the group, the +1 state becomes more stable. For example, Tl (+1) is more stable than Tl (+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 Tl (+1) is very stable.

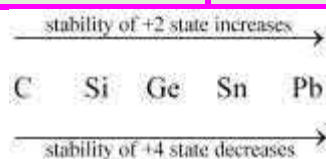
Group 13 element	Oxidation state
B	+3
Al	+3
Ga, In, Tl	+1, +3

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
C	+4
Si	+4
Ge, Sn, Pb	+2, +4



Question 11.2:

How can you explain higher stability of BCl_3 as compared to TlCl_3 ?

Answer 11.2:

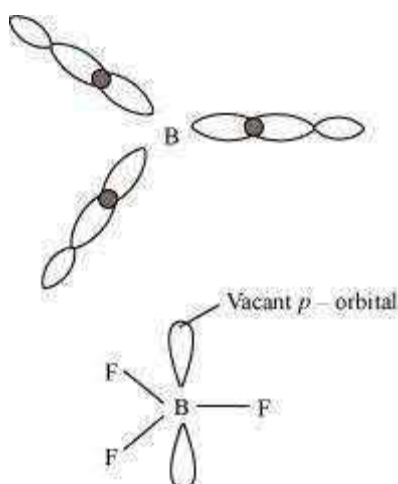
Boron and thallium belong to group 13 of the periodic table. In this group, the +1 oxidation state becomes more stable on moving down the group. BCl_3 is more stable than TlCl_3 because the +3 oxidation state of B is more stable than the +3 oxidation state of Tl. In Tl, the +3 state is highly oxidising and it reverts back to the more stable +1 state.

Question 11.3:

Why does boron trifluoride behave as a Lewis acid?

Answer 11.3:

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.



Question 11.4:

Consider the compounds, BCl₃ and CCl₄. How will they behave with water? Justify.

Answer 11.4:

Being a Lewis acid, BCl₃ readily undergoes hydrolysis. Boric acid is formed as a result.



CCl₄ completely resists hydrolysis. Carbon does not have any vacant orbital. Hence, it cannot accept electrons from water to form an intermediate. When CCl₄ and water are mixed, they form separate layers.

**Question 11.5:**

Is boric acid a protic acid? Explain.

Answer 11.5:

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



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

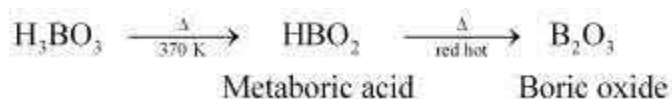
Question 11.6:

Explain what happens when boric acid is heated.

Answer 11.6:

On heating orthoboric acid (H₃BO₃) at 370 K or above, it changes to metaboric acid (HBO₂).

On further heating, this yields boric oxide B₂O₃.



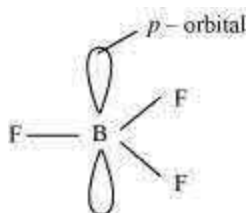
Question 11.7:

Describe the shapes of BF_3 and BH_4^- . Assign the hybridisation of boron in these species.

Answer 11.7:

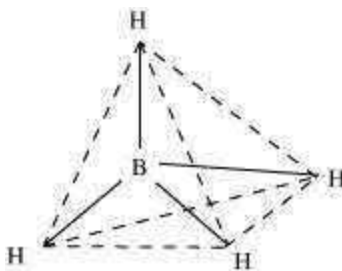
(i) BF_3

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 sp orbitals of three halogen atoms. Boron is sp^2 hybridised in BF_3 .



(ii) BH_4^-

Boron-hydride ion (BH_4^-) is formed by the sp^3 hybridisation of boron orbitals. Therefore, it is tetrahedral in structure.

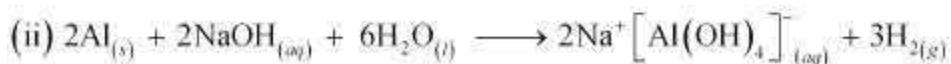
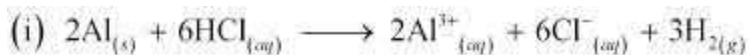


Question 11.8:

Write reactions to justify amphoteric nature of aluminium.

Answer 11.8:

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



Question 11.9:

What are electron deficient compounds? Are BCl_3 and $SiCl_4$ electron deficient species?

Explain.

Answer 11.9:

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) BCl_3

BCl_3 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) $SiCl_4$

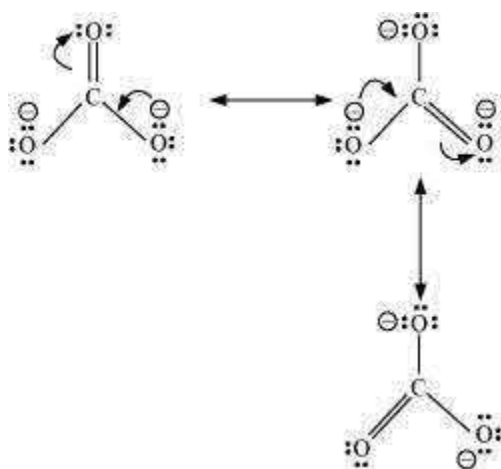
The electronic configuration of silicon is $ns^2 np^2$. This indicates that it has four valence electrons. After it forms four covalent bonds with four chlorine atoms, its electron count increases to eight. Thus, $SiCl_4$ is not an electron-deficient compound.

Question 11.10:

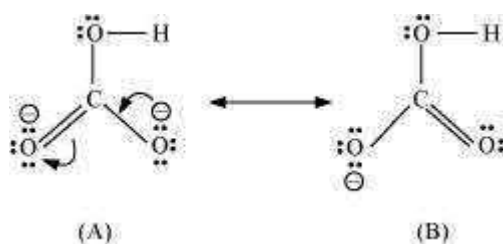
Write the resonance structures of CO_3^{2-} and HCO_3^{-} .

Answer 11.10:

(a) CO_3^{2-}



(b) HCO_3^-



There are only two resonating structures for the bicarbonate ion.

Question 11.11:

What is the state of hybridisation of carbon in (a) CO_3^{2-} (b) diamond (c) graphite?

Answer 11.11

The state of hybridisation of carbon in:

(a) CO_3^{2-}

C in CO_3^{2-} is sp^2 hybridised and is bonded to three oxygen atoms.

(b) Diamond

Each carbon in diamond is sp^3 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.

Question 11.12:

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

Answer 11.12:

Diamond	Graphite
It has a crystalline lattice.	It has a layered structure.
In diamond, each carbon atom is sp^3 hybridised and is bonded to four other carbon atoms through a σ bond.	In graphite, each carbon atom is sp^2 hybridised and is bonded to three other carbon atoms through a σ bond. The fourth electron forms a π 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.

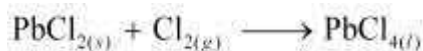
Question 11.13:

Rationalise the given statements and give chemical reactions:

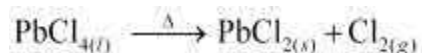
- Lead (II) chloride reacts with Cl_2 to give $PbCl_4$.
- Lead (IV) chloride is highly unstable towards heat.
- Lead is known not to form an iodide, PbI_4 .

Answer 11.13:

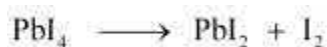
(a) Lead belongs to group 14 of the periodic table. The two oxidation states displayed by this group is +2 and +4. On moving down the group, the +2 oxidation state becomes more stable and the +4 oxidation state becomes less stable. This is because of the inert pair effect. Hence, $PbCl_4$ is much less stable than $PbCl_2$. However, the formation of $PbCl_4$ takes place when chlorine gas is bubbled through a saturated solution of $PbCl_2$.



(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).



(c) Lead is known not to form PbI_4 . 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^- to I_2 and itself gets reduced to Pb(II).

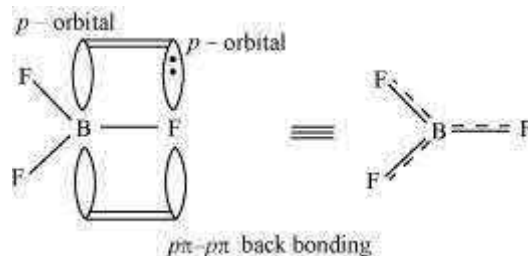


Question 11.14:

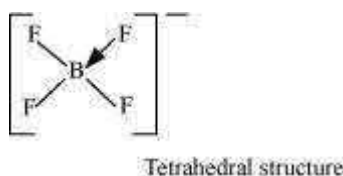
Suggest reasons why the B–F bond lengths in BF_3 (130 pm) and BF_4^- (143 pm) differ.

Answer 11.14:

The B–F bond length in BF_3 is shorter than the B–F bond length in BF_4^- . BF_3 is an electron-deficient species. With a vacant p -orbital on boron, the fluorine and boron atoms undergo $p\pi-p\pi$ back-bonding to remove this deficiency. This imparts a double bond character to the B–F bond.



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 σ bonds and the double-bond character is lost. This accounts for a B–F bond length of 143 pm in BF_4^- ion.

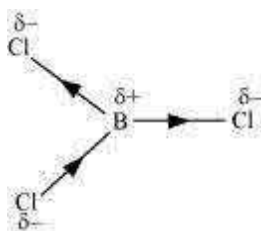


Question 11.15:

If B–Cl bond has a dipole moment, explain why BCl₃ molecule has zero dipole moment.

Answer 11.15:

As a result of the difference in the electronegativities of B and Cl, the B–Cl bond is polar in nature. However, the BCl₃ molecule is non-polar. This is because BCl₃ is trigonal planar in shape. It is a symmetrical molecule. Hence, the respective dipole-moments of the B–Cl bond cancel each other, thereby causing a zero-dipole moment.

**Question 11.16:**

Aluminium trifluoride is insoluble in anhydrous HF but dissolves on addition of NaF. Aluminium trifluoride precipitates out of the resulting solution when gaseous BF₃ is bubbled through. Give reasons.

Answer 11.16:

Hydrogen fluoride (HF) is a covalent compound and has a very strong intermolecular hydrogen-bonding. Thus, it does not provide ions and aluminium fluoride (AlF₃) does not dissolve in it. Sodium fluoride (NaF) is an ionic compound and when it is added to the mixture, AlF₃ dissolves. This is because of the availability of free F⁻. The reaction involved in the process is:



When boron trifluoride (BF₃) 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 BF₃ is added to the solution, B replaces Al from the complexes according to the following reaction:



Question 11.17:

Suggest a reason as to why CO is poisonous.

Answer 11.17:

Carbon monoxide is highly-poisonous because of its ability to form a complex with haemoglobin. The CO-Hb complex is more stable than the O₂-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 O₂-Hb complex.

Question 11.18:

How is excessive content of CO₂ responsible for global warming?

Answer 11.18:

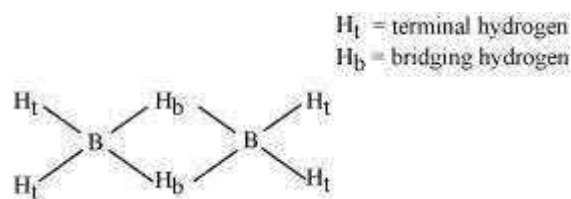
Carbon dioxide is a very essential gas for our survival. However, an increased content of CO₂ in the atmosphere poses a serious threat. An increment in the combustion of fossil fuels, decomposition of limestone, and a decrease in the number of trees has led to greater levels of carbon dioxide. Carbon 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.

Question 11.19:

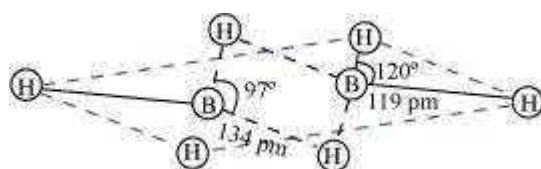
Explain structures of diborane and boric acid.

Answer 11.19:**(a) Diborane**

B₂H₆ is an electron-deficient compound. B₂H₆ has only 12 electrons – 6 e⁻ from 6 H atoms and 3 e⁻ each 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:

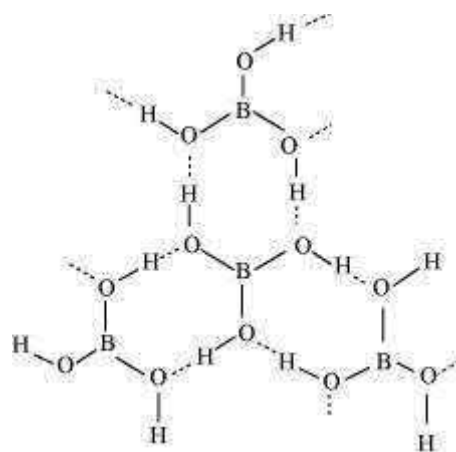


2 boron and 4 terminal hydrogen atoms (H_t) lie in one plane, while the other two bridging hydrogen atoms (H_b) lie in a plane perpendicular to the plane of boron atoms. Again, of the two bridging hydrogen atoms, one H atom lies above the plane and the other lies below the plane. The terminal bonds are 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.



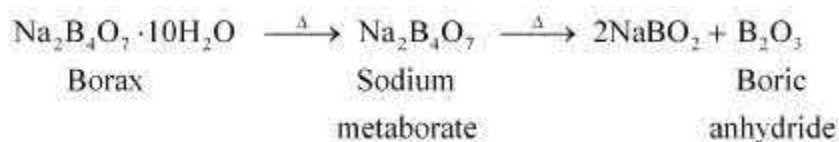
Question 11.20:

What happens when

- (a) Borax is heated strongly,
- (b) Boric acid is added to water,
- (c) Aluminium is treated with dilute NaOH,
- (d) BF₃ is reacted with ammonia?

Answer 11.20:

- (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.



- (b) When boric acid is added to water, it accepts electrons from ⁻OH ion.



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



- (d) BF₃ (a Lewis acid) reacts with NH₃ (a Lewis base) to form an adduct. This results in a complete octet around B in BF₃.

**Question 11.21:**

Explain the following reactions

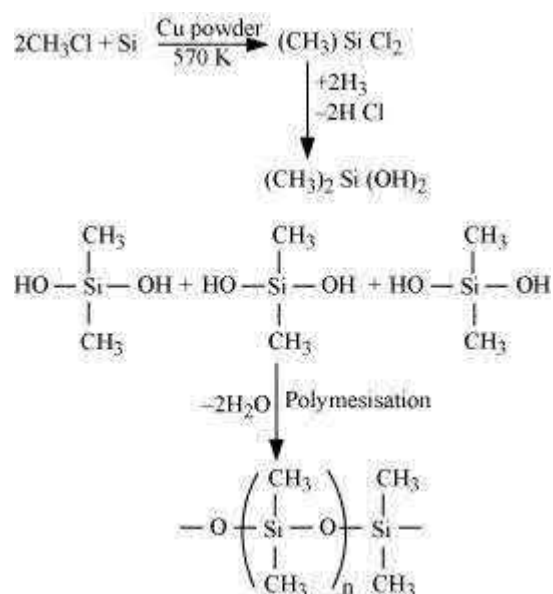
- (a) Silicon is heated with methyl chloride at high temperature in the presence of copper;
- (b) Silicon dioxide is treated with hydrogen fluoride;

(c) CO is heated with ZnO;

(d) Hydrated alumina is treated with aqueous NaOH solution.

Answer 11.21:

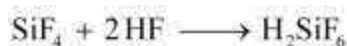
(a) When silicon reacts with methyl chloride in the presence of copper (catalyst) and at a temperature of about 537 K, a class of organosilicon polymers called methylsubstituted chlorosilanes (MeSiCl_3 , Me_2SiCl_2 , Me_3SiCl , and Me_4Si) are formed.



(b) When silicon dioxide (SiO_2) is heated with hydrogen fluoride (HF), it forms silicon tetrafluoride (SiF_4). 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.



The SiF_4 formed in this reaction can further react with HF to form hydrofluorosilicic acid.



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



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



Question 11.22:

Give reasons:

- (i) Conc. HNO_3 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 11.23:

(i) Concentrated HNO_3 can 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.



(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, Mg, 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.

Question 11.23:

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

Answer 11.23:

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.

Question 11.24:

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

Answer 11.24:

Atomic radius (in pm)	
Aluminium	143
Gallium	135

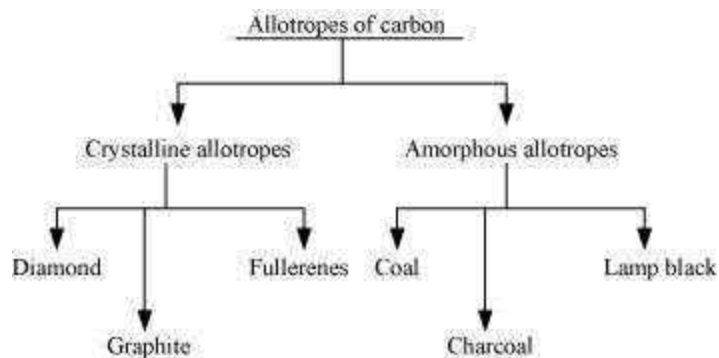
Although Ga has one shell more than Al, its size is lesser than Al. This is because of the poor shielding effect of the $3d$ -electrons. The shielding effect of d -electrons is very poor and the effective nuclear charge experienced by the valence electrons in gallium is much more than it is in the case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

Question 11.25:

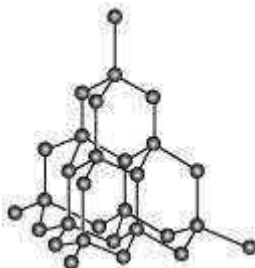
What are allotropes? Sketch the structure of two allotropes of carbon namely diamond and graphite. What is the impact of structure on physical properties of two allotropes?

Answer 11.25:

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.

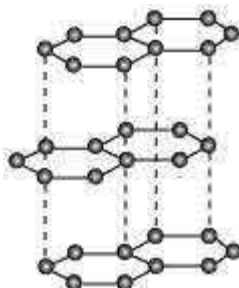


Diamond:



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^2 hybridised carbon, arranged in the form of layers. These layers are held together by weak van der Waals' forces. These layers can slide over each other, making graphite soft and slippery. Therefore, it is used as a lubricant.

Question 11.26:

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

CO, B_2O_3 , SiO_2 , CO_2 , Al_2O_3 , PbO_2 , Tl_2O_3

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

Answer 11.26:

(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.



(3) SiO_2 = Acidic

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



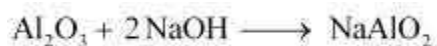
(4) CO_2 = Acidic

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



(5) Al_2O_3 = Amphoteric

Amphoteric substances react with both acids and bases. Al_2O_3 reacts with both NaOH and H_2SO_4 .



(6) PbO_2 = Amphoteric

Amphoteric substances react with both acids and bases. PbO_2 reacts with both NaOH and H_2SO_4 .



(7) Tl_2O_3 = Basic

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



Question 11.27:

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 11.27:

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 TlCl_3 and Tl_2O_3 . It resembles alkali metals in compounds Tl_2O and TlCl .

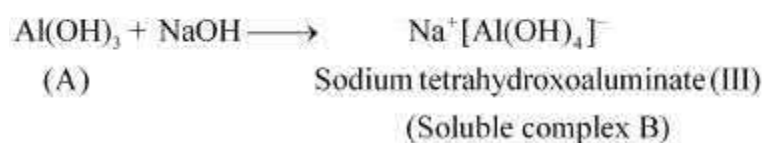
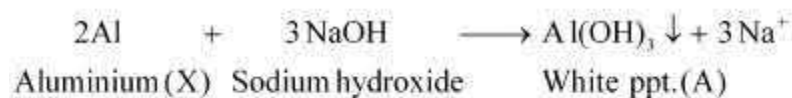
Question 11.28:

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.

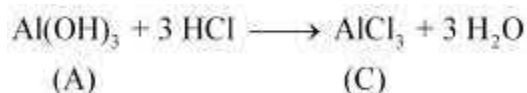
Answer 11.28:

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.

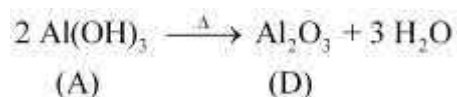
The white precipitate (compound A) obtained is aluminium hydroxide. The compound B formed when an excess of the base is added is sodium tetrahydroxoaluminate(III).



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



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.



Question 11.29:

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

Answer 11.29:

(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.

Question 11.30:

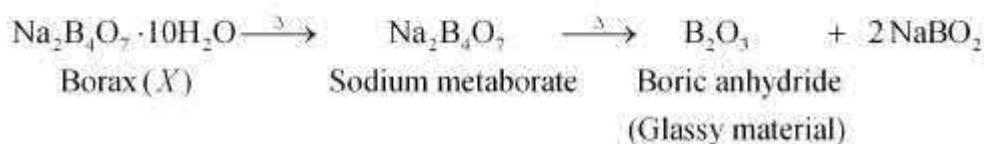
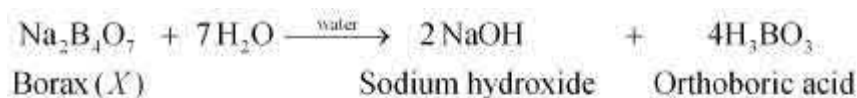
A certain salt X , gives the following results.

- (i) Its aqueous solution is alkaline to litmus.
- (ii) It swells up to a glassy material Y on strong heating.
- (iii) When conc. H_2SO_4 is added to a hot solution of X , white crystal of an acid Z separates out.

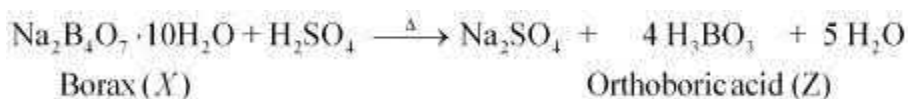
Write equations for all the above reactions and identify X , Y and Z .

Answer 11.30:

The given salt is alkaline to litmus. Therefore, X is a salt of a strong base and a weak acid. Also, when X is strongly heated, it swells to form substance Y . Therefore, X must 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, Y must be a mixture of sodium metaborate and boric anhydride.

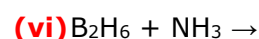
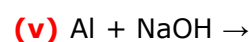
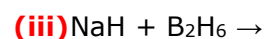
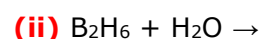
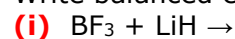


When concentrated acid is added to borax, white crystals of orthoboric acid (Z) are formed.

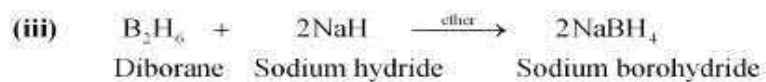
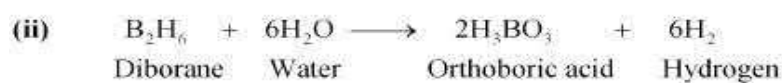
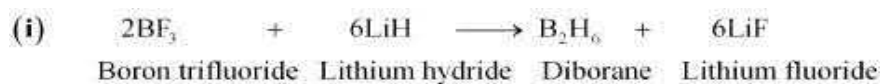


Question 11.31:

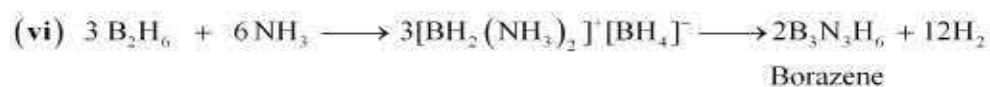
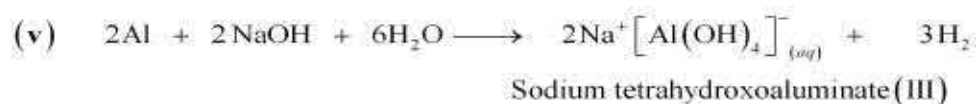
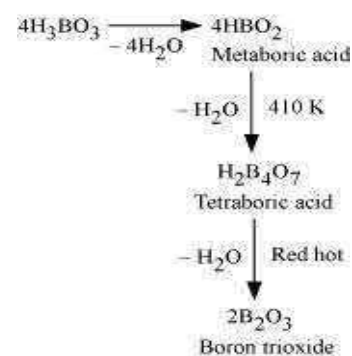
Write balanced equations for:



Answer 11.31:



(iv)

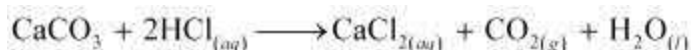


Question 11.32:

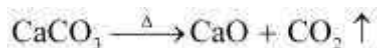
Give one method for industrial preparation and one for laboratory preparation of CO and CO₂ each.

Answer 11.32:**Carbon dioxide**

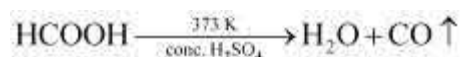
In the laboratory, CO₂ can be prepared by the action of dilute hydrochloric acid on calcium carbonate. The reaction involved is as follows:



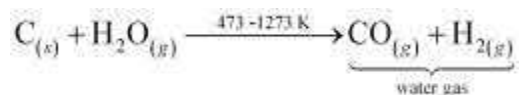
CO₂ is commercially prepared by heating limestone. The reaction involved is as follows:

**Carbon monoxide**

In the laboratory, CO is prepared by the dehydration of formic acid with conc. H₂SO₄, at 373 K. The reaction involved is as follows:



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

**Question 11.33:**

An aqueous solution of borax is

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

Answer 11.33:

(c) Borax is a salt of a strong base (NaOH) and a weak acid (H₃BO₃). It is, therefore, basic in nature.

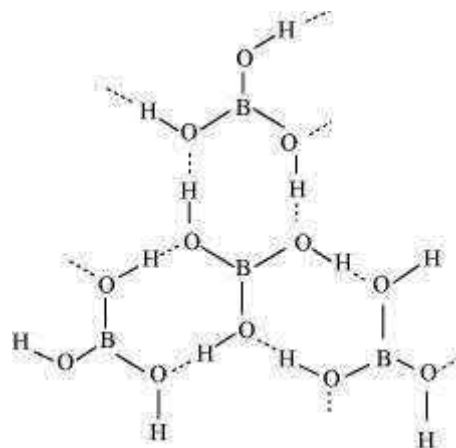
Question 11.34:

Boric acid is polymeric due to

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

Answer 11.34:

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



Question 11.35:

The type of hybridisation of boron in diborane is

- (a) sp (b) sp^2 (c) sp^3 (d) dsp^2

Answer 11.35:

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

Question 11.36:

Thermodynamically the most stable form of carbon is

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

Answer 11.36:

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

Question 11.37:

Elements of group 14

- (a) exhibit oxidation state of +4 only
(b) exhibit oxidation state of +2 and +4

(c) form M^{2-} and M^{4+} ion

(d) form M^{2+} and M^{4+} ions

Answer 11.37:

(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
C	+4
Si	+4
Ge, Sn, Pb	+2, +4

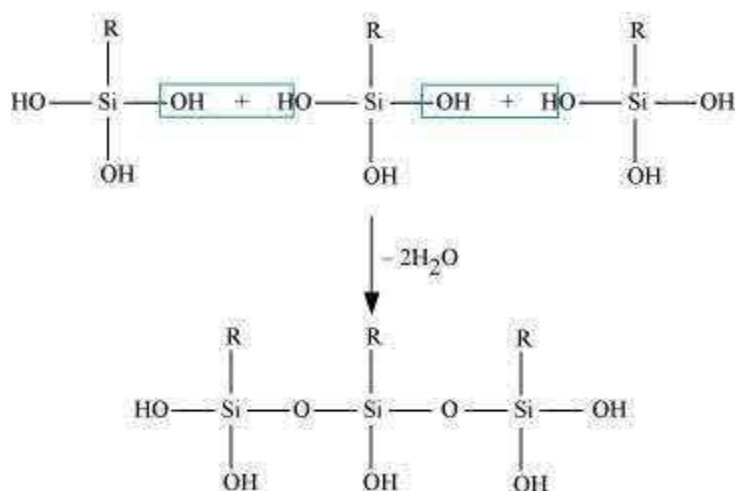
Question 11.38:

If the starting material for the manufacture of silicones is $RSiCl_3$, write the structure of the product formed.

Answer 11.38:



(i)



(ii)

