# (Chapter 7)(The p – Block Elements) XII

# **Intext Questions**

# Question 7.1:

Why are pentahalides more covalent than trihalides?

Answer

In pentahalides, the oxidation state is +5 and in trihalides, the oxidation state is +3. Since the metal ion with a high charge has more polarizing power, pentahalides are more covalent than trihalides.

## Question 7.2:

Why is BiH<sub>3</sub> the strongest reducing agent amongst all the hydrides of Group 15 elements?

Answer

As we move down a group, the atomic size increases and the stability of the hydrides of group 15 elements decreases. Since the stability of hydrides decreases on moving from  $NH_3$  to  $BiH_3$ , the reducing character of the hydrides increases on moving from  $NH_3$  to  $BiH_3$ .

## Question 7.3:

Why is N<sub>2</sub> less reactive at room temperature?

Answer

The two N atoms in  $N_2$  are bonded to each other by very strong triple covalent bonds. The bond dissociation energy of this bond is very high. As a result,  $N_2$  is less reactive at room temperature.

## Question 7.4:

Mention the conditions required to maximise the yield of ammonia.

Answer

Ammonia is prepared using the Haber's process. The yield of ammonia can be maximized under the following conditions: (i) High pressure ( $\sim$  200 atm)

(ii) A temperature of ~700 K

(iii) Use of a catalyst such as iron oxide mixed with small amounts of K<sub>2</sub>O and Al<sub>2</sub>O<sub>3</sub>

# Question 7.5:

How does ammonia react with a solution of Cu<sup>2+</sup>? Answer

NH<sub>3</sub> acts as a Lewis base. It donates its electron pair and forms a linkage with metal ion.

$$\operatorname{Cu}_{(aq)}^{2+} + 4\operatorname{NH}_{3(aq)} \leftrightarrow \left[\operatorname{Cu}(\operatorname{NH}_3)_4\right]_{(aq)}^{2+}$$

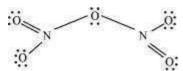
Blue

Deep blue

# Question 7.6:

What is the covalence of nitrogen in N<sub>2</sub>O<sub>5</sub>?

Answer



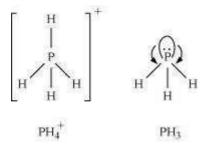
From the structure of  $N_2O_5$ , it is evident that the covalence of nitrogen is 4.

## Question 7.7:

Bond angle in  $\overset{PH_{4}^{+}}{}$  is higher than that in PH<sub>3</sub>. Why?

## Answer

In PH<sub>3</sub>, P is  $sp^3$  hybridized. Three orbitals are involved in bonding with three hydrogen atoms and the fourth one contains a lone pair. As lone pair-bond pair repulsion is stronger than bond pair-bond pair repulsion, the tetrahedral shape associated with  $sp^3$  bonding is changed to pyramidal. PH<sub>3</sub> combines with a proton to form  $PH_4^+$  in which the lone pair is absent. Due to the absence of lone pair in  $PH_4^+$ , there is no lone pair-bond pair repulsion. Hence, the bond angle in  $PH_4^+$  is higher than the bond angle in PH<sub>3</sub>.



# Question 7.8:

What happens when white phosphorus is heated with concentrated NaOH solution in an inert atmosphere of  $CO_2$ ?

Answer

White phosphorous dissolves in boiling NaOH solution (in a CO<sub>2</sub> atmosphere) to give phosphine, PH<sub>3</sub>.

$$P_4 + 3 \text{ NaOH} + 3 \text{ H}_2\text{O} \longrightarrow P\text{H}_3 + 3 \text{ NaH}_2\text{PO}_2$$

Phosphine Sodium hypophosphite

## Question 7.9:

What happens when PCI<sub>5</sub> is heated?

Answer

All the bonds that are present in  $PCl_5$  are not similar. It has three equatorial and two axial bonds. The equatorial bonds are stronger than the axial ones. Therefore, when  $PCl_5$  is heated strongly, it decomposes to form  $PCl_3$ .

$$PCl_5 \xrightarrow{heat} PCl_3 + Cl_2$$

## Question 7.10:

Write a balanced equation for the hydrolytic reaction of PCI<sub>5</sub> in heavy water. Answer

$$PCl_5 + D_2O \longrightarrow POCl_3 + 2DCl_2$$
  
 $POCl_3 + 3D_2O \longrightarrow D_3PO_4 + 3DCl_3$ 

Therefore, the net reaction can be written as

$$PCl_5 + 4D_2O \longrightarrow D_3PO_4 + 5DCl$$

# Question 7.11:

What is the basicity of H<sub>3</sub>PO<sub>4</sub>?

Answer

H<sub>3</sub>PO<sub>4</sub>

$$H_3PO_4 = P$$
 $HO OH$ 
 $OH$ 

Since there are three OH groups present in  $H_3PO_4$ , its basicity is three i.e., it is a tribasic acid.

## Question 7.12:

What happens when H<sub>3</sub>PO<sub>3</sub> is heated?

Answer

 $H_3PO_3$ , on heating, undergoes disproportionation reaction to form  $PH_3$  and  $H_3PO_4$ . The oxidation numbers of P in  $H_3PO_3$ ,  $PH_3$ , and  $H_3PO_4$  are +3, -3, and +5 respectively. As the oxidation number of the same element is decreasing and increasing during a particular reaction, the reaction is a disproportionation reaction.

$$4H_3PO_3 \xrightarrow{\Delta} 3H_3PO_4 + PH_3$$
Orthophosphorous acid Orthophosphoric acid Phosphine
(+3) (+5) (-3)

## Question 7.13:

List the important sources of sulphur.

Answer

Sulphur mainly exists in combined form in the earth's crust primarily as sulphates [gypsum (CaSO<sub>4</sub>.2H<sub>2</sub>O), Epsom salt (MgSO<sub>4</sub>.7H<sub>2</sub>O), baryte (BaSO<sub>4</sub>)] and sulphides [(galena (PbS), zinc blends (ZnS), copper pyrites (CuFeS<sub>2</sub>)].

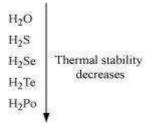
## Question 7.14:

Write the order of thermal stability of the hydrides of Group 16 elements.

#### Answer

The thermal stability of hydrides decreases on moving down the group. This is due to a decrease in the bond dissociation enthalpy (H-E) of hydrides on moving down the group.

## Therefore,



# Question 7.15:

Why is H<sub>2</sub>O a liquid and H<sub>2</sub>S a gas?

## Answer

 $H_2O$  has oxygen as the central atom. Oxygen has smaller size and higher electronegativity as compared to sulphur. Therefore, there is extensive hydrogen bonding in  $H_2O$ , which is absent in  $H_2S$ . Molecules of  $H_2S$  are held together only by weak van der Waal's forces of attraction.

Hence,  $H_2O$  exists as a liquid while  $H_2S$  as a solid.

# Question 7.16:

Which of the following does not react with oxygen directly?

Zn, Ti, Pt, Fe

## Answer

Pt is a noble metal and does not react very easily. All other elements, Zn, Ti, Fe, are quite reactive. Hence, oxygen does not react with platinum (Pt) directly.

# Question 7.17:

Complete the following reactions:

(i) 
$$C_2H_4 + O_2 \rightarrow$$

$$C_2H_4 + 3O_2 \longrightarrow 2CO_2 + 2H_2O$$
Ethene Oxygen Carbon dioxide Water
$$4Al + 3O_2 \longrightarrow 2Al_2O_3$$
Aluminium Oxygen Alumina
(i)

(ii)

## Question 7.18:

Why does O<sub>3</sub> act as a powerful oxidising agent?

Answer

Ozone is not a very stable compound under normal conditions and decomposes readily on heating to give a molecule of oxygen and nascent oxygen. Nascent oxygen, being a free radical, is very reactive.

$$O_3 \xrightarrow{\Delta} O_2 + [O]$$
  
Ozone Oxygen Nascent oxygen

Therefore, ozone acts as a powerful oxidising agent.

# Question 7.19:

How is O<sub>3</sub> estimated quantitatively?

Answer

Quantitatively, ozone can be estimated with the help of potassium iodide. When ozone is made to react with potassium iodide solution buffered with a borate buffer (pH 9.2), iodine is liberated. This liberated iodine can be titrated against a standard solution of sodium thiosulphate using starch as an indicator. The reactions involved in the process are given below.

## Question 7.20:

What happens when sulphur dioxide is passed through an aqueous solution of Fe(III) salt? Answer

 $SO_2$  acts as a reducing agent when passed through an aqueous solution containing Fe(III) salt. It reduces Fe(III) to Fe(II) i.e., ferric ions to ferrous ions.

$$2 \text{Fe}^{3+} + \text{SO}_2 + 2 \text{H}_2 \text{O} \longrightarrow 2 \text{Fe}^{2+} + \text{SO}_4^{2-} + 4 \text{H}^+$$

#### Question 7.21:

Comment on the nature of two S-O bonds formed in  $SO_2$  molecule. Are the two S-O bonds in this molecule equal?

## Answer

The electronic configuration of S is  $1s^2 2s^2 2p^6 3s^2 3p^4$ .

During the formation of  $SO_2$ , one electron from 3p orbital goes to the 3d orbital and S undergoes  $sp^2$  hybridization. Two of these orbitals form sigma bonds with two oxygen atoms and the third contains a lone pair. p-orbital and d-orbital contain an unpaired electron each. One of these electrons forms  $p\pi$ -  $p\pi$  bond with one oxygen atom and the other forms  $p\pi$ -  $d\pi$  bond with the other molecule. This is the reason  $SO_2$  has a bent structure. Also, it is a resonance hybrid of structures  $\mathbf{I}$  and  $\mathbf{II}$ .

$$\begin{bmatrix} \ddot{\mathbf{S}} & \ddot{\mathbf{S}} & \ddot{\mathbf{S}} \\ \mathbf{O} & \mathbf{I} \end{bmatrix} \equiv \ddot{\mathbf{S}} & \ddot{\mathbf{S}} & \mathbf{I} & \mathbf{$$

Both S-O bonds are equal in length (143 pm) and have a multiple bond character.

# Question 7.22:

How is the presence of SO<sub>2</sub> detected?

#### Answer

SO<sub>2</sub> is a colourless and pungent smelling gas.

It can be detected with the help of potassium permanganate solution. When  $SO_2$  is passed through an acidified potassium permanganate solution, it decolonizes the solution

as it reduces  $$MnO_4^-$ ions to $Mn^{2+}$ ions.}$ 

$$5SO_2 + 2MnO_4^- + 2H_2O \longrightarrow 5SO_4^{2-} + 4H^+ + 2Mn^{2+}$$

## Question 7.23:

Mention three areas in which H<sub>2</sub>SO<sub>4</sub> plays an important role.

#### Answer

Sulphuric acid is an important industrial chemical and is used for a lot of purposes. Some important uses of sulphuric acid are given below.

- (i) It is used in fertilizer industry. It is used to make various fertilizers such as ammonium sulphate and calcium super phosphate.
- (ii) It is used in the manufacture of pigments, paints, and detergents.
- (iii) It is used in the manufacture of storage batteries.

# Question 7.24:

Write the conditions to maximize the yield of H<sub>2</sub>SO<sub>4</sub> by Contact process.

### Answer

Manufacture of sulphuric acid by Contact process involves three steps.

- **1.** Burning of ores to form SO<sub>2</sub>
- **2.** Conversion of  $SO_2$  to  $SO_3$  by the reaction of the former with  $O_2$

 $(V_2O_5$  is used in this process as a catalyst.)

**3.** Absorption of  $SO_3$  in  $H_2SO_4$  to give oleum ( $H_2S_2O_7$ )

The key step in this process is the second step. In this step, two moles of gaseous reactants combine to give one mole of gaseous product. Also, this reaction is exothermic. Thus, in accordance with Le Chatelier's principle, to obtain the maximum amount of  $SO_3$  gas, temperature should be low and pressure should be high.

## **Question 7.25:**

Why is 
$$K_{a_2} \ll K_{a_1}$$
 for H<sub>2</sub>SO<sub>4</sub> in water?

$$H_2SO_{4(aq)} + H_2O_{(l)} \longrightarrow H_3O_{(aq)}^+ + HSO_{4(aq)}^-; \quad K_{a_1} > 10$$

$${\rm HSO}^-_{4(aq)} + {\rm H_2O}_{(l)} \longrightarrow {\rm H_3O}^+_{(aq)} + {\rm SO}^-_{4(aq)}; \qquad K_{a_2} = 1.2 \times 10^{-2}$$

$$K_{a_1} >> K_{a_2}$$

It can be noticed that  $K_{a_1} >> K_{a_2}$ 

This is because a neutral H<sub>2</sub>SO<sub>4</sub> has a much higher tendency to lose a proton than the negatively charged  ${\rm HSO_4^-}$ . Thus, the former is a much stronger acid than the latter.

## Question 7.26:

Considering the parameters such as bond dissociation enthalpy, electron gain enthalpy and hydration enthalpy, compare the oxidising power of F<sub>2</sub> and Cl<sub>2</sub>.

## Answer

Fluorine is a much stronger oxidizing agent than chlorine. The oxidizing power depends on three factors.

- 1. Bond dissociation energy
- 2. Electron gain enthalpy
- 3. Hydration enthalpy

The electron gain enthalpy of chlorine is more negative than that of fluorine. However, the bond dissociation energy of fluorine is much lesser than that of chlorine. Also, because of its small size, the hydration energy of fluorine is much higher than that of chlorine. Therefore, the latter two factors more than compensate for the less negative electron gain enthalpy of fluorine. Thus, fluorine is a much stronger oxidizing agent than chlorine.

## Question 7.27:

Give two examples to show the anomalous behaviour of fluorine.

Answer

## Anomalous behaviour of fluorine

- (i) It forms only one oxoacid as compared to other halogens that form a number of oxoacids.
- (ii) Ionisation enthalpy, electronegativity, and electrode potential of fluorine are much higher than expected.

## Question 7.28:

Sea is the greatest source of some halogens. Comment.

#### Answer

Sea water contains chlorides, bromides, and iodides of Na, K, Mg, and Ca. However, it primarily contains NaCl. The deposits of dried up sea beds contain sodium chloride and carnallite, KCl.MgCl<sub>2</sub>.6H<sub>2</sub>O. Marine life also contains iodine in their systems. For example, sea weeds contain upto 0.5% iodine as sodium iodide. Thus, sea is the greatest source of halogens.

## Question 7.29:

Give the reason for bleaching action of Cl<sub>2</sub>.

#### Answer

When chlorine reacts with water, it produces nascent oxygen. This nascent oxygen then combines with the coloured substances present in the organic matter to oxide them into colourless substances.

$$Cl_2 + H_2O \longrightarrow 2HCl + [O]$$

Coloured substances + [O]  $\rightarrow$  Oxidized colourless substance

## Question 7.30:

Name two poisonous gases which can be prepared from chlorine gas.

#### Answer

Two poisonous gases that can be prepared from chlorine gas are

- (i) Phosgene (COCl<sub>2</sub>)
- (ii) Mustard gas (CICH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CI)

Question 7.31:

Why is ICl more reactive than  $I_2$ ?

Answer

ICl is more reactive than I<sub>2</sub> because I-Cl bond in ICl is weaker than I-I bond in I<sub>2</sub>.

Question 7.32:

Why is helium used in diving apparatus?

Answer

Air contains a large amount of nitrogen and the solubility of gases in liquids increases with increase in pressure. When sea divers dive deep into the sea, large amount of nitrogen dissolves in their blood. When they come back to the surface, solubility of nitrogen decreases and it separates from the blood and forms small air bubbles. This leads to a dangerous medical condition called bends. Therefore, air in oxygen cylinders used for diving is diluted with helium gas. This is done as He is sparingly less soluble in blood.

Question 7.33:

Balance the following equation:  $XeF_6 + H_2O \rightarrow XeO_2F_2 + HF$ 

Answer

**Balanced equation** 

 $XeF_6 + 2 H_2O \rightarrow XeO_2F_2 + 4 HF$ 

Question 7.34:

Why has it been difficult to study the chemistry of radon?

Answer

It is difficult to study the chemistry of radon because it is a radioactive substance having a half-life of only 3.82 days. Also, compounds of radon such as  $RnF_2$  have not been isolated. They have only been identified.

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