# The p-Block Elements

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Chapter	1110	P-DIOCK	LICITICITIS	LACIDISC	Jointhorn

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Q1:

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.

#### Q2:

Why is BiH3the 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<sub>3</sub>to BiH<sub>3</sub>, the reducing character of the hydrides increases on moving from NH<sub>3</sub>to BiH<sub>3</sub>.

### Q3:

Why is N₂less 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.

### Q4:

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 (∝½200 atm)

- (ii) A temperature of ∝1/4700 K
- (iii) Use of a catalyst such as iron oxide mixed with small amounts of K2O and Al2O3

### Q5:

How does ammonia react with a solution of Cu2+?

#### Answer:

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

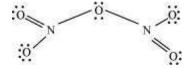
$$Cu^{2+}_{(aq)} + 4NH_{3(aq)} \leftrightarrow \left[Cu(NH_3)_4\right]^{2+}_{(aq)}$$

Blue Deep blue

#### Q6:

What is the covalence of nitrogen in N₂O₅?

#### Answer:



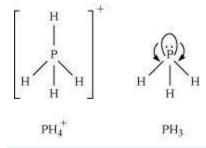
From the structure of N<sub>2</sub>O<sub>5</sub>, it is evident that the covalence of nitrogen is 4.

### Q7:

Bond angle in  $${{PH}_{4}^{+}}$$  is higher than that in PH3. 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>.



Q8:

What happens when white phosphorus is heated with concentrated NaOHsolution in an inert atmosphere of CO<sub>2</sub>?

### Answer:

White phosphorous dissolves in boiling NaOH solution (in a CO2atmosphere) to give phosphine, PH3.

$$P_4 + 3 \text{ NaOH} + 3 \text{ H}_2 \text{O} \longrightarrow PH_3 + 3 \text{ NaH}_2 PO_2$$

Phosphine Sodium hypophosphite

Q9:

What happens when PCI₅is heated?

### Answer:

All the bonds that are present in PCI<sub>5</sub> are not similar. It has three equatorial and two axial bonds. The equatorial bonds are stronger than the axial ones. Therefore, when PCI<sub>5</sub> is heated strongly, it decomposes to form PCI<sub>3</sub>.

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

Q10:

Write a balanced equation for the hydrolytic reaction of PCI₅in 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

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

### Q11:

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

### Answer:

 $H_3PO_4$ 

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

Since there are three OH groups present in H<sub>3</sub>PO<sub>4</sub>, its basicity is three i.e., it is a tribasic acid.

### Q12:

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)

### Q13:

List the important sources of sulphur.

### Answer:

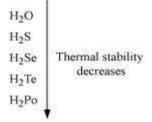
Sulphur mainly exists in combined form in the earth's crust primarily as sulphates [gypsum (CaSO<sub>4</sub>.2H<sub>2</sub>O), Epsom salt (MgSO<sub>4</sub>.7H<sub>2</sub>O), baryte (BaSO<sub>4</sub>)] and sulphides [(galena (PbS), zinc blends (ZnS), copper pyrites (CuFeS<sub>2</sub>)].

# Q14:

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

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,



#### Q15:

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<sub>2</sub>O exists as a liquid while H<sub>2</sub>S as a gas.

### Q16:

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.

### Q17:

Complete the following reactions:

(i)  $C_2H_4 + O_2 \rightarrow$ 

(ii) 4AI +  $3O_2 \rightarrow$ 

$$C_2H_4 + 3O_2 \longrightarrow 2CO_2 + 2H_2O$$
(i) Ethene Oxygen Carbon dioxide Water

 $4Al + 3O_2 \longrightarrow 2Al_2O_3$ 
(ii) Aluminium Oxygen Alumina

#### Q18:

Why does O₃act 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.

### Q19:

How is O₃estimated 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.

### Q20:

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

 $SO_2$  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.

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

#### Q21:

Comment on the nature of two S-O bonds formed in SO<sub>2</sub>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<sub>2</sub>, 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  $p\tilde{A}\hat{a},\neg$   $p\tilde{A}\hat{a},\neg$  bond with one oxygen atom and the other forms $p\tilde{A}\hat{a},\neg$  da0 bond with the other oxygen. This is the reason SO<sub>2</sub> has a bent structure. Also, it is a resonance hybrid of structures I and II.

$$\begin{bmatrix} \ddot{\mathbf{S}} & \ddot{\mathbf{S}} & \ddot{\mathbf{S}} \\ \mathbf{O} & \ddot{\mathbf{S}} & \mathbf{O} \end{bmatrix} \equiv \ddot{\mathbf{S}} & \overset{\mathsf{143 pm}}{\mathsf{109.5}^{\circ}} \\ \mathbf{O} & \mathbf{O} & \mathbf{O} & \mathbf{O} \\ \mathbf{O} & \mathbf{O} \\ \mathbf{O} & \mathbf{O} & \mathbf{O} \\ \mathbf{O} \\ \mathbf{O} & \mathbf{O} \\ \mathbf{O} & \mathbf{O} \\ \mathbf{O} & \mathbf{O} \\ \mathbf{O} \\ \mathbf{O} & \mathbf{O} \\ \mathbf{O} \\ \mathbf{O} & \mathbf{O} \\ \mathbf$$

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

# Q22:

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.

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

#### Q23:

Mention three areas in which H₂SO₄plays an important role.

Sulphuric acid is an important industrial chemicaland 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.

#### Q24:

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<sub>2</sub>to SO<sub>3</sub>by the reaction of the former with O<sub>2</sub>

(V₂O₅is used in this process as a catalyst.)

3. Absorption of SO<sub>3</sub>in H<sub>2</sub>SO<sub>4</sub>to give oleum (H<sub>2</sub>S<sub>2</sub>O<sub>7</sub>)

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 SO₃gas, temperature should be low and pressure should be high.

Q25:

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

#### Answer:

$$\begin{split} & \text{H}_2\text{SO}_{4(aq)} + \text{H}_2\text{O}_{(l)} \longrightarrow \text{H}_3\text{O}_{(aq)}^+ + \text{HSO}_{4(aq)}^-; \quad K_{a_1} > 10 \\ & \text{HSO}_{4(aq)}^- + \text{H}_2\text{O}_{(l)} \longrightarrow \text{H}_3\text{O}_{(aq)}^+ + \text{SO}_{4(aq)}^-; \qquad K_{a_2} = 1.2 \times 10^{-2} \end{split}$$
 It can be noticed that  $K_{a_1} >> K_{a_2}$ 

This is because a neutral H₂SO₄has a much higher tendency to lose a proton than the negatively charged Thus, the former is a much stronger acid than the latter.

### Q26:

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

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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.

#### Q27:

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.

### Q28:

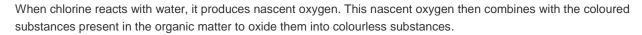
Sea is the greatest source of some halogens. Comment.

#### 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.

### Q29:

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



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

Coloured substances + [O] â†' Oxidized colourless substance

#### Q30:

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 (COCI<sub>2</sub>)
- (ii) Mustard gas (CICH<sub>2</sub>CH<sub>2</sub>SCH<sub>2</sub>CH<sub>2</sub>CI)

#### Q31:

Why is ICI more reactive than I<sub>2</sub>?

## Answer:

ICI is more reactive than I2because I-CI bond in ICI is weaker than I-I bond in I2.

#### Q32:

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.

### Q33:

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

### **Balanced equation**

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

#### Q34:

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<sub>2</sub> have not been isolated. They have only been identified.

Exercise: Solutions of Questions on Page Number: 207

Q1:

Discuss the general characteristics of Group 15 elements with reference to their electronic configuration, oxidation state, atomic size, ionisation enthalpy and electronegativity.

### Answer:

# General trends in group 15 elements

- (i) **Electronic configuration**: All the elements in group 15 have 5 valence electrons. Their general electronic configuration is  $ns^2 np^3$ .
- (ii) Oxidation states: All these elements have 5 valence electrons and require three more electrons to complete their octets. However, gaining electrons is very difficult as the nucleus will have to attract three more electrons. This can take place only with nitrogen as it is the smallest in size and the distance between the nucleus and the valence shell is relatively small. The remaining elements of this group show a formal oxidation state of -3 in their covalent compounds. In addition to the -3 state, N and P also show -1 and -2 oxidation states.

All the elements present in this group show +3 and +5 oxidation states. However, the stability of +5 oxidation state decreases down a group, whereas the stability of +3 oxidation state increases. This happens because of the inert pair effect.

#### (iii) Ionization energy and electronegativity

First ionization decreases on moving down a group. This is because of increasing atomic sizes. As we move down a group, electronegativity decreases, owing to an increase in size.

(iv) Atomic size: On moving down a group, the atomic size increases. This increase in the atomic size is attributed to an increase in the number of shells.

Q2:

### Why does the reactivity of nitrogen differ from phosphorus?

#### Answer:

Nitrogen is chemically less reactive. This is because of the high stability of its molecule,  $N_2$ . In  $N_2$ , the two nitrogen atoms form a triple bond. This triple bond has very high bond strength, which is very difficult to break. It is because of nitrogen's small size that it is able to form pÃâ,¬-pÃâ,¬bonds with itself. This property is not exhibited by atoms such as phosphorus. Thus, phosphorus is more reactive than nitrogen.

#### Q3:

Discuss the trends in chemical reactivity of group 15 elements.

#### Answer:

General trends in chemical properties of group - 15

### (i) Reactivity towards hydrogen:

The elements of group 15 react with hydrogen to form hydrides of type  $EH_3$ , where E = N, P, As, Sb, or Bi. The stability of hydrides decreases on moving down from  $NH_3$  to  $BiH_3$ .

## (ii) Reactivity towards oxygen:

The elements of group 15 form two types of oxides:  $E_2O_3$  and  $E_2O_5$ , where E = N, P, As, Sb, or Bi. The oxide with the element in the higher oxidation state is more acidic than the other. However, the acidic character decreases on moving down a group.

#### (iii) Reactivity towards halogens:

The group 15 elements react with halogens to form two series of salts:  $EX_3$  and  $EX_5$ . However, nitrogen does not form  $NX_5$  as it lacks the d-orbital. All trihalides (except  $NX_3$ ) are stable.

### (iv) Reactivity towards metals:

The group 15 elements react with metals to form binary compounds in which metals exhibit -3 oxidation states.

#### Q4:

Why does NH<sub>3</sub>form hydrogen bond but PH<sub>3</sub>does not?

#### Answer:

Nitrogen is highly electronegative as compared to phosphorus. This causes a greater attraction of electrons towards nitrogen in  $NH_3$ than towards phosphorus in  $PH_3$ . Hence, the extent of hydrogen bonding in  $PH_3$  is very less as compared to  $NH_3$ .

### Q5:

How is nitrogen prepared in the laboratory? Write the chemical equations of the reactions involved.

Answer:

An aqueous solution of ammonium chloride is treated with sodium nitrite.

$$NH_4Cl_{(aq)} + NaNO_{2(aq)} \longrightarrow N_{2(g)} + 2H_2O_{(l)} + NaCl_{(aq)}$$

NO and HNO₃ are produced in small amounts. These are impurities that can be removed on passing nitrogen gas through aqueous sulphuric acid, containing potassium dichromate.

Q6:

How is ammonia manufactured industrially?

Answer:

Ammonia is prepared on a large-scale by the Haber's process.

 $N_2(g) + 3H_2(g) \tilde{A} \phi \hat{a} \in \hat{a} \in 2NH_3(g)$ 

Q7:

Illustrate how copper metal can give different products on reaction with HNO<sub>3</sub>.

Answer:

Concentrated nitric acid is a strong oxidizing agent. It is used for oxidizing most metals. The products of oxidation depend on the concentration of the acid, temperature, and also on the material undergoing oxidation.

$$3Cu + 8HNO_{3(dilute)} \longrightarrow 3Cu(NO_3)_2 + 2NO + 4H_2O$$

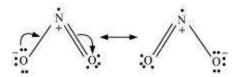
$$Cu + 4HNO_{3(conc.)} \longrightarrow Cu(NO_3)_2 + 2NO_2 + 2H_2O$$

Q8:

Give the resonating structures of NO<sub>2</sub> and N<sub>2</sub>O<sub>5</sub>.

Answer:

(1)



(2)

#### Q9:

The HNH angle value is higher than HPH, HAsH and HSbH angles. Why? [Hint: Can be explained on the basis of *sp*³hybridisation in NH₃and only *s-p* bonding between hydrogen and other elements of the group].

#### Answer:

Hydride NH<sub>3</sub> PH<sub>3</sub> AsH<sub>3</sub> SbH<sub>3</sub>

H-M-H angle 107° 92° 91° 90°

The above trend in the H-M-H bond angle can be explained on the basis of the electronegativity of the central atom. Since nitrogen is highly electronegative, there is high electron density around nitrogen. This causes greater repulsion between the electron pairs around nitrogen, resulting in maximum bond angle. We know that electronegativity decreases on moving down a group. Consequently, the repulsive interactions between the electron pairs decrease, thereby decreasing the H-M-H bond angle.

# Q10:

Why does  $R_3P=0$  exist but  $R_3N=0$  does not (R = alkyl group)?

## Answer:

N(unlike P) lacks the d-orbital. This restricts nitrogen to expand its coordination number beyond four. Hence, R<sub>3</sub>N=O does not exist.

### Q11:

Explain why NH<sub>3</sub>is basic while BiH<sub>3</sub>is only feebly basic.

### Answer:

# NH<sub>3</sub>is distinctly basic while BiH<sub>3</sub>is feebly basic.

Nitrogen has a small size due to which the lone pair of electrons is concentrated in a small region. This means that the charge density per unit volume is high. On moving down a group, the size of the central atom increases and the charge gets distributed over a large area decreasing the electron density. Hence, the electron donating capacity of group 15 element hydrides decreases on moving down the group.

### Q12:

Nitrogen exists as diatomic molecule and phosphorus as P4. Why?

# Answer:

Nitrogen owing to its small size has a tendency to form  $p\tilde{A}\hat{a},\neg-p\tilde{A}\hat{a},\neg$ multiple bonds with itself. Nitrogen thus forms a very stable diatomic molecule,  $N_2$ . On moving down a group, the tendency to form  $p\tilde{A}\hat{a},\neg-p\tilde{A}\hat{a},\neg$ bonds decreases (because of the large size of heavier elements). Therefore, phosphorus (like other heavier metals) exists in the  $P_4$ state.

#### Q13:

Write main differences between the properties of white phosphorus and red phosphorus.

### Answer:

White phosphorus	Red Phosphorus			
It is a soft and waxy solid. It possesses a garlic smell.	It is a hard and crystalline solid, without any smell.			
It is poisonous.	It is non-poisonous.			
It is insoluble in water but soluble in carbon disulphide.	It is insoluble in both water and carbon disulphide.			
It undergoes spontaneous combustion in air.	It is relatively less reactive.			
In both solid and vapour states, it exists as a P <sub>4</sub> molecule.	It exists as a chain of tetrahedral P <sub>4</sub> units.  P P P P P P P P P P P P P P P P P P			

### Q14:

Why does nitrogen show catenation properties less than phosphorus?

Catenation is much more common in phosphorous compounds than in nitrogen compounds. This is because of the relative weakness of the N-N single bond as compared to the P-P single bond. Since nitrogen atom is smaller, there is greater repulsion of electron density of two nitrogen atoms, thereby weakening the N-N single bond.

#### Q15:

Give the disproportionation reaction of H<sub>3</sub>PO<sub>3</sub>.

#### Answer:

On heating, orthophosphorus acid  $(H_3PO_3)$  disproportionates to give orthophosphoric acid  $(H_3PO_4)$  and phosphine  $(PH_3)$ . The oxidation states of P in various species involved in the reaction are mentioned below.

$$4H_3 \stackrel{+3}{P}O_3 \longrightarrow 3H_3 \stackrel{+5}{P}O_4 + \stackrel{-3}{P}H_3$$

#### Q16:

Can PCI₅act as an oxidising as well as a reducing agent? Justify.

#### Answer:

PCl₅can only act as an oxidizing agent. The highest oxidation state that P can show is +5. In PCl₅, phosphorus is in its highest oxidation state (+5). However, it can decrease its oxidation state and act as an oxidizing agent.

### Q17:

Justify the placement of O, S, Se, Te and Po in the same group of the periodic table in terms of electronic configuration, oxidation state and hydride formation.

#### Answer:

The elements of group 16 are collectively called chalcogens.

(i) Elements of group 16 have six valence electrons each. The general electronic configuration of these elements is  $ns^2np^4$ , where nv arises from 2 to 6.

### (ii) Oxidation state:

Asthese elements have six valence electrons ( $ns^2$   $np^4$ ), they should display an oxidation state of -2. However, only oxygen predominantly shows the oxidation state of -2 owing to its high electronegativity. It also exhibits the oxidation state of -1 ( $H_2O_2$ ), zero ( $O_2$ ), and +2 ( $OF_2$ ). However, the stability of the -2 oxidation state decreases on moving down a group due to a decrease in the electronegativity of the elements. The heavier elements of the group show an oxidation state of +2, +4, and +6 due to the availability of d-orbitals.

#### (iii) Formation of hydrides:

These elements form hydrides of formula  $H_2E$ , where E = O, S, Se, Te, PO. Oxygen and sulphur also form hydrides of type  $H_2E_2$ . These hydrides are quite volatile in nature.

#### Q18:

Why is dioxygen a gas but sulphur a solid?

#### Answer:

Oxygen is smaller in size as compared to sulphur. Due to its smaller size, it can effectively form  $p\tilde{A}\hat{a},\neg p\tilde{A}\hat{a},\neg p$ 

#### Q19:

Knowing the electron gain enthalpy values for O  $\rightarrow$  O and O  $\rightarrow$  O as -141 and 702 kJ mol espectively, how can you account for the formation of a large number of oxides having O as and not O?

(Hint: Consider lattice energy factor in the formation of compounds).

#### Answer:

Stability of an ionic compound depends on its lattice energy. More the lattice energy of a compound, more stable it will be.

Lattice energy is directly proportional to the charge carried by an ion. When a metal combines with oxygen, the lattice energy of the oxide involving O²-ion is much more than the oxide involving Oʻ-ion. Hence, the oxide having O²-ions are more stable than oxides having Oʻ-. Hence, we can say that formation of O²-is energetically more favourable than formation of Oʻ-.

#### Q20:

Which aerosols deplete ozone?

#### Answer:

Freons or chlorofluorocarbons (CFCs) are aerosols that accelerate the depletion of ozone. In the presence of ultraviolet radiations, molecules of CFCs break down to form chlorine-free radicals that combine with ozone to form oxygen.

#### Q21:

Describe the manufacture of H<sub>2</sub>SO<sub>4</sub>by contact process?

#### Answer:

Sulphuric acid is manufactured by the contact process. It involves the following steps:

### Step (i):

Sulphur or sulphide ores are burnt in air to form SO<sub>2</sub>.

### Step (ii):

By a reaction with oxygen, SO₂is converted into SO₃in the presence of V₂O₅as a catalyst.

$$2\mathrm{SO}_{2(g)} + \mathrm{O}_{2(g)} \xrightarrow{\quad v_2\mathrm{O}_5 \quad} 2\mathrm{SO}_{3(g)}$$

# Step (iii):

SO<sub>3</sub>produced is absorbed on H<sub>2</sub>SO<sub>4</sub>to give H<sub>2</sub>S<sub>2</sub>O<sub>7</sub>(oleum).

$$SO_3 + H_2SO_4 \longrightarrow H_2S_2O_7$$

This oleum is then diluted to obtain H<sub>2</sub>SO<sub>4</sub>of the desired concentration.

In practice, the plant is operated at 2 bar (pressure) and 720 K (temperature). The sulphuric acid thus obtained is 96-98% pure.

### Q22:

How is SO₂an air pollutant?

### Answer:

Sulphur dioxide causes harm to the environment in many ways:

- **1.** It combines with water vapour present in the atmosphere to form sulphuric acid. This causes acid rain. Acid rain damages soil, plants, and buildings, especially those made of marble.
- **2.** Even in very low concentrations, SO<sub>2</sub> causes irritation in the respiratory tract. It causes throat and eye irritation and can also affect the larynx to cause breathlessness.
- **3.** It is extremely harmful to plants. Plants exposed to sulphur dioxide for a long time lose colour from their leaves. This condition is known as chlorosis. This happens because the formation of chlorophyll is affected by the presence of sulphur dioxide.

### Q23:

Why are halogens strong oxidising agents?

The general electronic configuration of halogens is $np^5$ , where $n=2-6$ . Thus, halogens need only one more electron to complete their octet and to attain the stable noble gas configuration. Also, halogens are highly electronegative with low dissociation energies and high negative electron gain enthalpies. Therefore, they have a high tendency to gain an electron. Hence, they act as strong oxidizing agents.
Q24 :
Explain why fluorine forms only one oxoacid, HOF.
Answer:
Fluorine forms only one oxoacid i.e., HOF because of its high electronegativity and small size.
Q25 :
Explain why inspite of nearly the same electronegativity, oxygen forms hydrogen bonding while chlorine does not.
Answer:
Both chlorine and oxygen have almost the same electronegativity values, but chlorine rarely forms hydrogen bonding. This is because in comparison to chlorine, oxygen has a smaller size and as a result, a higher electron density per unit volume.
Q26 :
Write two uses of CIO <sub>2</sub> .
Answer:
Uses of CIO <sub>2</sub> :
(i) It is used for purifying water.
(ii) It is used as a bleaching agent.
Q27 :
Why are halogens coloured?

Almost all halogens are coloured. This is because halogens absorb radiations in the visible region. This results in the excitation of valence electrons to a higher energy region. Since the amount of energy required for excitation differs for each halogen, each halogen displays a different colour.

### Q28:

Write the reactions of F2and Cl2with water.

### Answer:

$$Cl_2 + H_2O \longrightarrow HCl + HOCl$$

(i) Hydrochloric acid Hypochlorous acid

(ii) 
$$2F_{2(g)} + 2H_2O_{(I)} \longrightarrow 4H^+_{(aq)} + 4F^-_{(aq)} + O_{2(g)} + 4HF_{(aq)}$$

#### Q29:

How can you prepare Cl<sub>2</sub> from HCl and HCl from Cl<sub>2</sub>? Write reactions only.

#### Answer:

(i) Cl<sub>2</sub>can be prepared from HCl by Deacon's process.

$$4\,\mathrm{HCl} + \mathrm{O_2} \xrightarrow{\quad \mathrm{CuCl_2} \quad} 2\,\mathrm{Cl_2} + 2\,\mathrm{H_2O}$$

(ii) HCl can be prepared from Cl2 on treating it with water.

$$Cl_2 + H_2O \longrightarrow HCl + HOCl$$

Hydrochloric acid Hypochlorous acid

# Q30:

What inspired N. Bartlett for carrying out reaction between Xe and PtF6?

### Answer:

Neil Bartlett initially carried out a reaction between oxygen and PtF<sub>6</sub>. This resulted in the formation of a red compound,  $O_2^+[PtF_6^-]^-$ .

Later, he realized that the first ionization energy of oxygen (1175 kJ/mol) and Xe (1170 kJ/mol) is almost the same. Thus, he tried to prepare a compound with Xe and PtF<sub>6</sub>. He was successful and a red-coloured  $Xe^+\Gamma PtF^-I^-$ 

compound, 
$$Xe^+[PtF_6^-]^-$$
 was formed.

#### Q31:

What are the oxidation states of phosphorus in the following:

- (i) H<sub>3</sub>PO<sub>3</sub> (ii) PCI<sub>3</sub> (iii) Ca<sub>3</sub>P<sub>2</sub>
- (iv) Na<sub>3</sub>PO<sub>4</sub> (v) POF<sub>3</sub>?

Let the oxidation state of p be x

(i) H<sub>3</sub>PO<sub>3</sub>

$$3+x+3(-2)=0$$

$$3 + x - 6 = 0$$

$$x - 3 = 0$$

$$x = +3$$

(ii) PCl<sub>3</sub>

$$x+3(-1)=0$$

$$x - 3 = 0$$

$$x = +3$$

(iii) Ca<sub>3</sub>P<sub>2</sub>

$$3(+2)+2(x)=0$$

$$6 + 2x = 0$$

$$2x = -6$$

$$x = -3$$

(iv) Na<sub>3</sub>PO<sub>4</sub>

$$3(+1) + x + 4(-2) = 0$$

$$3 + x - 8 = 0$$

$$x - 5 = 0$$

$$x = +5$$

(v) POF<sub>3</sub>

$$x + (-2) + 3(-1) = 0$$

$$x - 5 = 0$$

$$x = +5$$

# Q32:

Write balanced equations for the following:

- (i) NaCl is heated with sulphuric acid in the presence of MnO<sub>2</sub>.
- (ii) Chlorine gas is passed into a solution of Nal in water.

(i) 
$$4\text{NaCl} + \text{MnO}_2 + 4\text{H}_2\text{SO}_4 \longrightarrow \text{MnCl}_2 + 4\text{NaHSO}_4 + 2\text{H}_2\text{O} + \text{Cl}_2$$

(ii) 
$$Cl_2 + NaI \longrightarrow 2 NaCl + I_2$$

#### Q33:

How are xenon fluorides XeF<sub>2</sub>, XeF<sub>4</sub>and XeF<sub>6</sub>obtained?

#### Answer:

XeF<sub>2</sub>, XeF<sub>4</sub>,and XeF<sub>6</sub> are obtained by a direct reaction between Xe and F2. The condition under which the reaction is carried out determines the product.

$$Xe_{(g)} + F_{2(g)} \xrightarrow{-673K,1bar} XeF_{2(s)}$$
(Excess)

 $Xe_{(g)} + 2F_{2(g)} \xrightarrow{-873K,7bar} XeF_{4(s)}$ 
(1:5 ratio)

 $Xe_{(g)} + 3F_{2(g)} \xrightarrow{-573K,60-70bar} XeF_{6(s)}$ 
(1:20 ratio)

#### Q34:

With what neutral molecule is CIO isoelectronic? Is that molecule a Lewis base?

#### Answer:

CIO is isoelectronic to CIF. Also, both species contain 26 electrons in all as shown.

Total electrons CIO = 17 + 8 + 1 = 26

In CIF = 
$$17 + 9 = 26$$

CIF acts like a Lewis base as it accepts electrons from F to form CIF<sub>3</sub>.

### Q35:

How are XeO₃and XeOF₄prepared?

(i) XeO₃can be prepared in two ways as shown.

$$6XeF_4 + 12H_2O \longrightarrow 4Xe + 2XeO_3 + 24HF + 3O_2$$

$$XeF_6 + 3H_2O \longrightarrow XeO_3 + 6HF$$

(ii) XeOF<sub>4</sub>can be prepared using XeF<sub>6</sub>.

$$XeF_6 + H_2O \longrightarrow XeOF_4 + 2HF$$

#### Q36:

Arrange the following in the order of property indicated for each set:

- (i) F<sub>2</sub>, Cl<sub>2</sub>, Br<sub>2</sub>, l<sub>2</sub>- increasing bond dissociation enthalpy.
- (ii) HF, HCI, HBr, HI increasing acid strength.
- (iii) NH<sub>3</sub>, PH<sub>3</sub>, AsH<sub>3</sub>, SbH<sub>3</sub>, BiH<sub>3</sub>- increasing base strength.

#### Answer:

(i) Bond dissociation energy usually decreases on moving down a group as the atomic size increases. However, the bond dissociation energy of F2 is lower than that of Cl2 and Br2. This is due to the small atomic size of fluorine. Thus, the increasing order for bond dissociation energy among halogens is as follows:

$$I_2 < F_2 < Br_2 < Cl_2$$

(ii) HF < HCl < HBr < HI

The bond dissociation energy of H-X molecules where X = F, CI, Br, I, decreases with an increase in the atomic size. Since H-I bond is the weakest, HI is the strongest acid.

(iii) 
$$BiH_3 \tilde{A} c$$
"  $\hat{A} x SbH_3 < AsH_3 < PH_3 < NH_3$ 

On moving from nitrogen to bismuth, the size of the atom increases while the electron density on the atom decreases. Thus, the basic strength decreases.

### Q37:

Which one of the following does not exist?

- (i) XeOF<sub>4</sub> (ii) NeF<sub>2</sub>
- (iii) XeF<sub>2</sub> (iv) XeF<sub>6</sub>

# Answer:

NeF<sub>2</sub> does not exist.

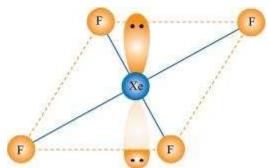
Give the formula and describe the structure of a noble gas species which is isostructural with:

- (i) ICl<sub>4</sub>
- (ii)  $IBr_2^-$
- (iii)  $BrO_3^-$

# Answer:

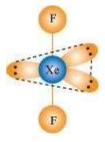
(i)

 ${\rm XeF_4} {\rm is}$  isoelectronic with  ${\rm \ }^{\rm \ } {\rm \ }^{\rm \ } {\rm \ }^{\rm \ }$  and has square planar geometry.



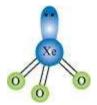
(ii)

 $\text{XeF}_{\text{\tiny 2}}\text{is}$  isoelectronic to  ${}^{1}\!Br_{2}^{-}$  and has a linear structure.



(iii)

 $\text{XeO}_{\text{s}}\text{is}$  isostructural to  ${\color{red}BrO_{3}^{-}}$  and has a pyramidal molecular structure.



# Q39:

Why do noble gases have comparatively large atomic sizes?

Noble gases do not form molecules. In case of noble gases, the atomic radii corresponds to van der Waal's radii. On the other hand, the atomic radii of other elements correspond to their covalent radii. By definition, van der Waal's radii are larger than covalent radii. It is for this reason that noble gases are very large in size as compared to other atoms belonging to the same period.

### Q40:

List the uses of Neon and argon gases.

#### Answer:

# Uses of neon gas:

- (i) It is mixed with helium to protect electrical equipments from high voltage.
- (ii) It is filled in discharge tubes with characteristic colours.
- (iii) It is used in beacon lights.

# Uses of Argon gas:

- (i) Argon along with nitrogen is used in gas-filled electric lamps. This is because Ar is more inert than N.
- (ii) It is usually used to provide an inert temperature in a high metallurgical process.
- (iii) It is also used in laboratories to handle air-sensitive substances.