# NCERT Solutions for Class 12 Chemistry Chapter 7 The p-Block Elements

# NCERT IN-TEXT QUESTIONS

Question 1.

Why are pentahalides more covalent than trihalides in the members of the nitrogen family? Answer:

The electronic configuration of the elements of nitrogen family (group 15) is  $ns^2p^3$ . Because of the inert pair effect, the valence s-electrons cannot be released easily for the bond formation. This means that the elements can form trivalent cation ( $E^{3+}$ ) by releasing valence p-electrons while it is difficult to form pentavalent cation ( $E^{5+}$ ). Under the circumstances, if all the five valence electrons are to be involved in the bond formation, the compounds showing pentavalency or +5 oxidation state must be of covalent nature. This is particularly the case in the higher members (Sb, Bi) of the family where the inert pair effect is quite prominent.

# Question 2.

Why is  $\text{BiH}_{\scriptscriptstyle 3}$  the strongest reducing agent amongst all the hydrides of Group 15 elements? Answer:

This is because as we move down the group, the size increases, as a result, length of E-H bond increases and its strength decreases, so that the bond can be broken easily to release  $H_2$  gas. Hence,  $BiH_3$  is the strongest reducing agent.

Question 3.

Why is  $N_2$  less reactive at room temperature?

Answer:

In the nitrogen molecule  $(N_2)$ , two atoms of nitrogen are linked by triple bond (N = N). Due to small atomic size of the element (atomic radius = 70

pm), the bond dissociation enthalpy is very high (946 kJ mol<sup>1</sup>). This means that it is quite difficult to cleave or break the triple bond at room temperature. As a result,  $N_2$  is less reactive at room temperature.

Question 4.

Mention the conditions required for the maximum yield of ammonia.

Answer:

In Haber's process, ammonia is formed by the following reaction.

 $N_2(g) + 3H_2(g)$ 

 $\frac{\text{Iron oxide (catalyst)}}{2NH_3} (g), \Delta_f H^\circ = -46.1 \text{ kJ mol}^{-1}$ 

According to Le-chatelier's principle, the favourable conditions for the maximum yield of ammonia are :

(i) Low temperature : But optimum temperature of 700 K is necessary to keep the forward reaction in progress.

(ii) High pressure : Pressure to the extent of about 200 atm is required.

(iii) Catalyst & promoter : In order to achieve the early attainment of equilibrium, iron oxide acts as catalyst. Along with that;  $K_2O$ ,  $Al_2O_3$  or Mo metal may act as the promoter to increase the efficiency of the catalyst.

## Question 5.

How does ammonia react with blue solution having Cu2+ ions?

Answer:

When ammonia gas is passed through blue solution containing Cu<sup>2+</sup> ions to give a soluble complex with deep blue colour.

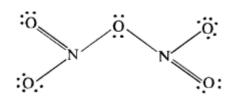
 $\operatorname{Cu}^{2+}(aq) + 4\operatorname{NH}_{4}\operatorname{OH}(aq) \longrightarrow [\operatorname{Cu}(\operatorname{NH}_{3})_{4}]^{2+}_{(aq)} + 4\operatorname{H}_{2}\operatorname{O}$ Tetraammine copper (II) ion.

(Deep blue solution)

Question 6.

What is the covalency of nitrogen in  $N_2O_5$ ? Answer:

The covalent structure of nitrogen pentoxide ( $N_2O_5$ ) is given. Since the nitrogen atom has shared four electron pairs, its covalency is four in the molecule of  $N_2O_5$ .



Question 7.

Bond angle in  $PH_{4^+}$  is higher than that in  $PH_{3^-}$ . Why?

Answer:

P in PH<sub>3</sub> is sp<sup>3</sup>-hybridized with 3 bond pairs and one lone pair around P. Due to stronger lp-bp repulsions than bp-bp repulsions, tetrahedral angle decreases from 109°28' to 93.6°. As a result, PH<sub>3</sub> is pyramidal. In PH<sub>4</sub>+, there are 4 bp's and no lone pair. As a result, there are only identical bp-bp repulsions so that PH<sub>4</sub>+ assumes tetrahedral geometry and the bond angle is 109°28'.Hence, bond angle of PH<sub>4</sub>+ > bond angle of PH<sub>3</sub>

Question 8.

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

Answer:

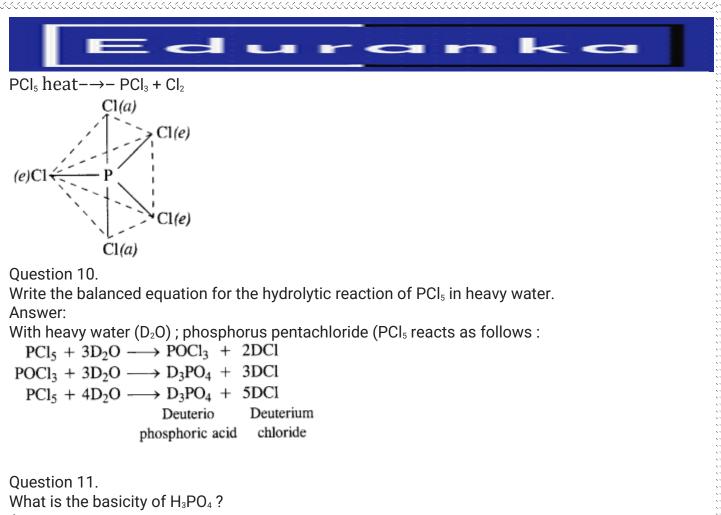
A mixture of sodium hypophosphite and phosphine gas is formed upon heating the reaction mixture in an inert atmosphere.

 $\begin{array}{c} P_4 + 3NaOH + 3H_2O \xrightarrow[(CO_2 \text{ atmosphere})]{Heat} & 3NaH_2PO_2 + PH_3\\ \text{White phosphorus} & Sod. hypo & Phosphine\\ \text{phosphite} & \end{array}$ 

Question 9. What happens when PCI₅ is heated?

Answer:

Upon heating,  $PCI_5$  dissociates to give molecules of  $PCI_3$  and  $CI_2$ . Actually, the two P-CI (a) bonds with more bond length break away from the molecule leaving three P-CI(e) bonds attached to the central P atom since these are more firmly linked



Answer:

The acid is tribasic since it has three P-OH bonds which can release  $H^+$  ions.

Question 12.

What happens when phosphorus acid ( $H_3PO_3$ ) is heated ? (C.B.S.E. 2008) Answer:

In phosphorus acid ( $H_3PO_3$ ), central atom P is in +3 oxidation state. Upon heating, it gives a mixture PH  $_3$  (P in -3 oxidation state) and  $H_3PO_4$  (P in +5 oxidation state). This means that phosphorus acid undergoes disproportionation reaction.

 $4H_3^{+3}PO_3 \xrightarrow{heat} PH_3^{-3} + 3H_3^{+5}PO_4$ Phosphine Phosphoric acid

Question 13.

List the important sources of sulphur.

Answer:

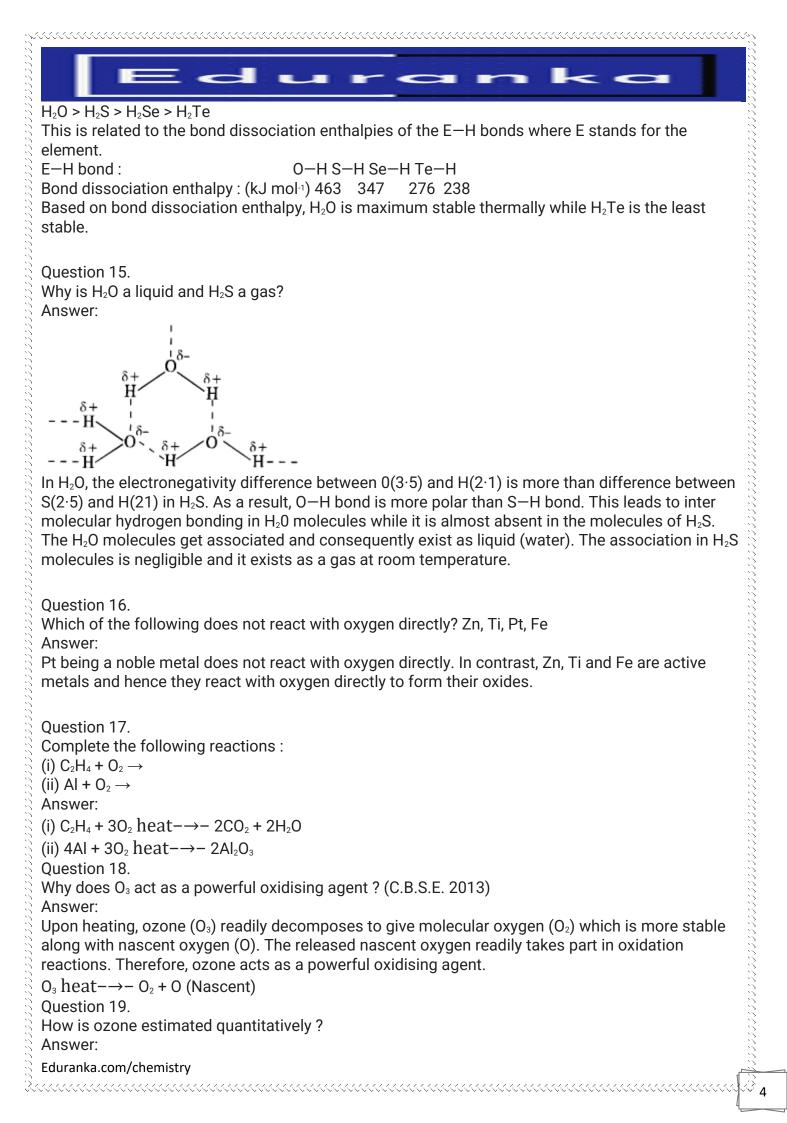
Combined sulphur exists as sulphates, such as gypsum, epsom, baryte and sulphides such as galena, zinc blende, copper pyrites, etc. Traces of sulphur occur as hydrogen sulphide in volcanoes. Few organic materials like eggs, proteins, garlic, onion, mustard, hair and wool contain sulphur. 0.03 – 0.1% sulphur is present in the earth's crust.

Question 14.

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

Answer:

The order of thermal stability of hydride is :





When ozone reacts with an excess of KI solution buffered with a borate buffer (pH = 9.2), iodine is liberated which can be titrated against standard solution of sodium thiosulphate. This is used as a method of estimation of ozone quantitatively.

Question 20.

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

The gas is a reducing agent and reduces a Fe(III) salt to Fe(II) salt.

 $SO_2 + 2H_2O \longrightarrow SO_4^{2-} + 4H^+ + 2e^-$ 2Fe<sup>3+</sup> + 2e<sup>-</sup>  $\longrightarrow$  2Fe<sup>2+</sup>

 $2Fe^{3+} + SO_2 + 2H_2O \longrightarrow 2Fe^{2+} + SO_4^{2-} + 4H^+.$ 

For example, ferric sulphate is reduced to ferrous sulphate.

 $Fe_2(SO_4)_3 + SO_2 + 2H_2O \longrightarrow 2FeSO_4 + 2H_2SO_4.$ 

Question 21.

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

Answer:

The two S-O bonds in the molecule are equal with bond length equal to 143 pm. This means that SO<sub>2</sub> molecule exhibits two resonating structures as shown below.

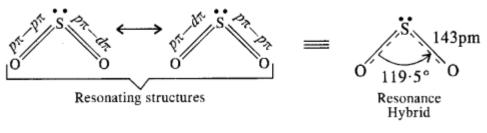


Figure 7.21. Structure of SO<sub>2</sub> (g)

Question 22. How is presence of  $SO_2$  detected ? Ans.

Presence of  $SO_2$  is detected by bringing a paper dipped in acidified potassium dichromate near the gas. If the paper turns green, it shows the presence of  $SO_2$  gas.

 $\begin{array}{c} K_2 Cr_2 O_7 + 4H_2 SO_4 & \longrightarrow K_2 SO_4 + Cr_2 (SO_4)_3 + 4H_2 O + 3[O] \\ \hline [SO_2 + 2H_2 O & \longrightarrow H_2 SO_4 + 2H] \times 3 \\ \hline 2H + O & \longrightarrow H_2 O] \times 3 \\ \hline K_2 Cr_2 O_7 + 3SO_2 + H_2 SO_4 & \longrightarrow K_2 SO_4 + Cr_2 (SO_4)_3 + H_2 O \\ & (Green) \end{array}$ 

Question 23.

Mention three areas in which  $H_2SO_4$  plays an important role.

Answer:

(i) It is used in the manufacture of fertilizers such as  $(NH_4)^2 SO_4$ , calcium superphosphate.



(ii)It is used as an electrolyte in storage batteries.

(iii)It is used in petroleum refining, detergent industry and in the manufacture of paints, pigments and dyes.

Question 24.

Write the conditions to maximise the yield of sulphuric acid by Contact process. Answer:

Catalytic oxidation of sulphur dioxide into sulphur trioxide. Sulphur dixocide is oxidised to sulphur trioxide with air in the presence of  $V_2O_5$  or platinised asbestos acting as the catalyst

 $2SO_2 + O_2 \xrightarrow{\text{catalyst}} 2SO_3$ ;  $\Delta H = -196.6 \text{ kJ mol}^{-1}$ 

Question 25.

Why is  $K_{a2} < K_{a1}$  for  $H_2SO_4$  in water ?

Answer:

 $H_2SO_4$  is a very strong acid in water largely because of its first ionisation to  $H_3O^+$  and  $HSO_4^-$  The ionisation of  $HSO_4^-$  to  $H_3O^+$  and  $SO_{4^{2-}}$  is very very small. That is why,  $Ka_2 < Ka_1$ .

## Question 26.

Considering the parameters such as bond dissociation enthalpy, electron gain enthalpy and hydration enthalpy, compare the oxidising power of  $F_2$  and  $Cl_2$ .

#### Answer:

Fluorine is a better oxidising agent than chlorine because  $E^{\circ}_{F2/F_{-}}$  is higher than  $E^{\circ}_{C12/C1_{-}}$  It is mainly due to low bond dissociation energy, high hydration energy and lower electron gain enthalpy, non-availability of d-orbitals in valence shell, that results in higher reduction potential of  $F_2$  than chlorine.

## Question 27.

Give two examples to show the anomalous behaviour of fluorine.

Answer:

The anomalous behaviour of fluorine, the first member of the halogen family as compared to the rest of the members is due to its very small size, very high electronegativity and absence of vacant d-orbitals in the valence shell. It is supported by the following points.

(i) Fluorine shows only negative oxidation state of -1 in its compounds. The other members exhibit both positive and negative oxidation states.

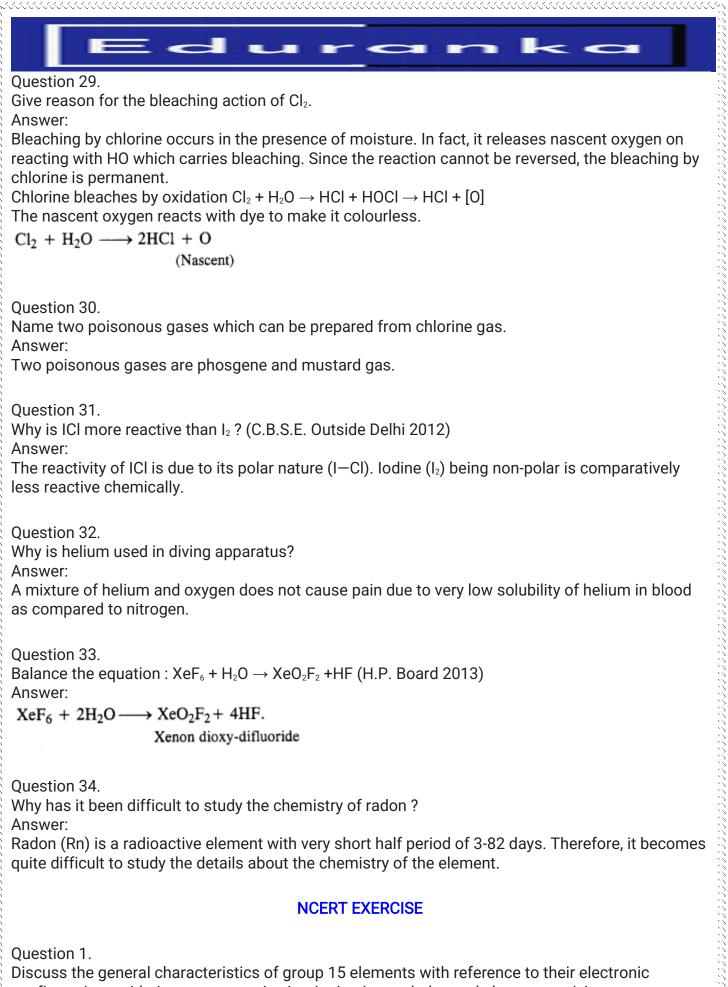
(ii) Fluorine forms hexafluoride with sulphur (SF $_6$ ). No other member of the family forms hexahalide with sulphur.

Question 28.

Sea is the greatest source of some halogens. Comment.

Answer:

The name halogen is a Greek Word meaning Isea salt forming'. Sea is a major source of a members of halogens particularly chlorine, bromine and iodine and they exist as the soluble salts of sodium, potassium, calcium, magnesium etc. The deposits of dried up sea water contain sodium chloride and carnallite (KCIMgCl<sub>2</sub>.6H<sub>2</sub>O). Sea weeds contain nearly 0-5 percent of iodine. Similarly Chile saltpeter contains about 0-2% of sodium iodate (NaIO<sub>3</sub>).



configuration, oxidation state, atomic size, ionisation enthalpy and electronegativity. Answer:



(i) Electronic configuration: All the elements in group 15 have 5 valence electrons. Their general electronic configuration is  $ns^2np^3$ 

## (ii) Oxidation states:

All these elements have 5 valence electrons and require three more electrons to complete 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 valance 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 the +5 oxidation state decreases down the group, whereas the stability of+3 oxidation state increases. This happens because of the inert pair effect.

(iii) Ionization energy and electronegativity First ionization energy 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.

Question 2.

Why does the reactivity of nitrogen differ from phosphorus?

Answer:

 $N_2$  exist as a diatomic molecule containing triple bond between two N-atoms. Due to the presence —of the triple bond between p the two N-atoms, the bond dissociation energy is large (941 .4 kJ mol<sup>-1</sup>). As a result of this  $N_2$  is inert and unreactive whereas, phosphorus exists as a tetratomic molecule, containing P – P single bond. Due to the presence of single bond, the bond dissociation energy is weaker (213 kJmol<sup>-1</sup>) than N a N triple bond (941 .4 kJ mol<sup>-1</sup>) and moreover due to presence of angular strain in P<sub>4</sub> tetrahedra. As a result of this, phosphorus is much more reactive than nitrogen.

Question 3.

Discuss the trends in chemical reactivity of group 15 elements. Answer:

(a) Hydrides: The hybrids are covalent with pyramidal structures <sup>(MH<sub>3</sub>)</sup> and the central atom is sp<sup>3</sup> hybrilised. The presence of the lone pair of electrons on the central atom distorts the geometry of the molecules and the bond angle less than that of a regular tetrahedron.

NH <sub>3</sub>	$PH_3$		AsH <sub>3</sub>	SbH <sub>3</sub>	BiH <sub>3</sub>
107·8°	93.6°		91.8°	91·3°	90°
An h	-	-			

**(b)** Halides: Elements of group 15 form two types of halides viz. trihalides and pentahalides. The halides are predominantly basic (Lewis bases) in nature and have lone pair of electrons (central atom is sp3 hybridized). The pentahalides are thermally less stable than the trihalides.



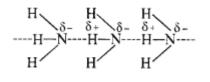
(c) Oxides: All the elements of this group form two types of oxides ie.,  $M_2O_3$  and  $M_2O_5$  and are called trioxides and pentoxides

Question 4.

Why does  $NH_3$  form hydrogen bonding while  $PH_3$  does not?

Answer:

The N–H bond in ammonia is quite polar on account of the electronegativity difference of N (3.0) and H (2.1). On the contrary, P–H bond in phosphine is almost non-polar because both P and H atoms have almost same electronegativity (21). Due to polarity, intermolecular hydrogen bonding is present in the molecules of ammonia but not in those of phosphine.



Question 5.

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

Answer:

Laboratory preparation: Dinitrogen is prepared in the laboratory by heating a solution containing an equivalent amount of sodium nitrite and ammonium chloride.

 $\begin{array}{rcl} \mathrm{NH_4Cl+\ NaNO_2} & \xrightarrow{\mathrm{Heat}} & \mathrm{NH_4NO_2 +\ NaCl} \\ & & \mathrm{Amm.\ nitrite} \\ & & \mathrm{NH_4NO_2} \xrightarrow{\mathrm{Heat}} & \mathrm{N_2} & + & \mathrm{2H_2O} \end{array}$ 

Question 6.

How is ammonia manufactured industrially?

Answer:

Ammonia is prepared on a commercial scale by Haber's Process from dinitrogen and dihydrogen by the following chemical reaction

 $N_2 + 3H_2 \rightleftharpoons 2NH_3$ ;  $\Delta_f H^\circ = -46.1 \text{ kJ mon}^{-1}$ 

Dihydrogen needed for the commercial preparation of ammonia is obtained by the electrolysis of water while dinitrogen is obtained from the liquefied air as a fractional distillation. The two gases are purified and also dried.

These are compressed to about 200 atmosphere pressure and are then led into the catalyst

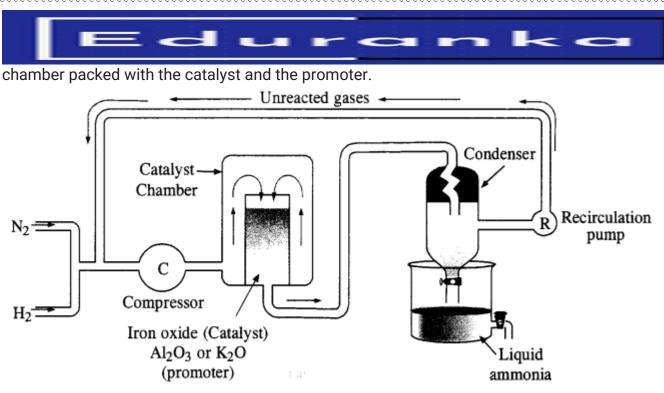


Figure 7.4. Haber's process for the manufacture of ammonia.

## Question 7.

Illustrate how copper metal gives different products on reaction with  $HNO_3$ . Answer:

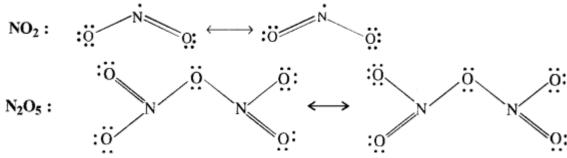
Anhydrous nitric acid is colourless, fuming, and pungent-smelling liquid. However, it acquires a yellowish-brown colour in the presence of sunlight. HNO<sub>3</sub> decomposes to give NO<sub>2</sub> gas which dissolves and imparts its yellowish-brown colour.

 $4HNO_3 \xrightarrow{Sun \ light} 4NO_2 + 2H_2O + O_2$ 

On strong heating, the acid decomposes to give  $NO_2$  and  $O_2$ 

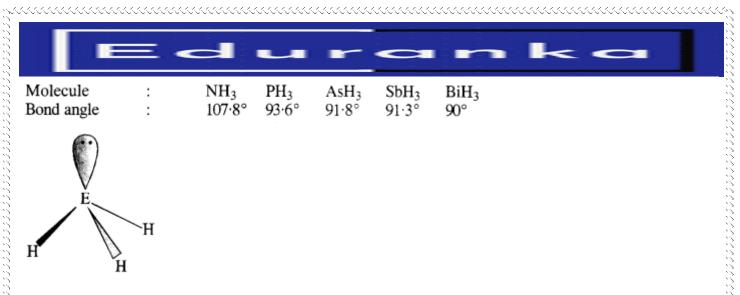
$$4\text{HNO}_3 \xrightarrow{\text{rieal}} 2\text{H}_2\text{O} + 4\text{NO}_2 + \text{O}_2$$

Question 8. Give the resonating structures of  $NO_2$  and  $N_2O_5$ . Answer:



Question 9.

The HNH angle value is higher than those of HPH, HAsH, and HSbH angles; why? Answer:



The above trend in the H-M-H bond angle can be explained on the basis of the electronegativity tif 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 HMH bond angle.

Question 10.

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

Answer:

Nitrogen does not contain d-orbitals. As a result, it cannot expand its covalency beyond four and cannot form  $p\pi - d\pi$  multiple bonds. In contrast, P contains the d-orbitals, and can expand its covalency beyond 4 and can form  $p\pi$ - $d\pi$  multiple bonds. Hence R<sub>3</sub>P = 0 exist but R<sub>3</sub>N = 0 does not.

Question 11.

Explain why is  $NH_3$  basic while  $PH_3$  is feebly basic in nature. (C.B.S.E. Outside Delhi 2008, 2009, Jharkhand Board 2009)

Answer:

NH3 is distinctly basic while  $\text{BiH}_{\mbox{\tiny 3}}$  is feebly basic.

Nitrogen has a small size due to which the lone pair of electrons are 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 elements hydrides decreases on moving down the group.

Question 12.

Nitrogen exists as a diatomic molecule ( $N_2$ ) while phosphorus a tetra-atomic molecule ( $P_4$ ). Why? Answer:

Nitrogen is diatomic gaseous molecule at ordinary temperature due to its ability to form  $p\pi - p\pi$  multiple bonds. The molecule has one  $\sigma$  and two  $\pi$  – bonds. Phosphorus exists as discrete tetratomic tetrahedral molecules as these are not capable of forming multiple bonds due to repulsion between non-bonded electrons of the inner core.

Question 13.

Write the main difference between the properties of white and red phosphorus. (C.B.S.E. Delhi 2012)

Property	White phosphorus	Red phosphorus	
1. Physical state	Soft waxy solid	Brittle powder	
2. Colour	White when freshly cut and becomes yellow on standing	Red in colour	
3. Odour	Garlic odour	Odourless	
<ol> <li>Physiological action</li> </ol>	Poisonous	Non-poisonous	
5. Solubility in water	Insoluble	Insoluble	
6. Solubility in CS <sub>2</sub>	Soluble	Insoluble	
7. Chemical reactivity	Very reactive	Comparitively less reactive	
8. Stability	Unstable	Stable in nature	
9. Molecular state	Exists as P <sub>4</sub> molecules	Exists in polymeric form	
10. Ignition temperature	Low (303 K)	High (543 K)	
11. Phosphorescence	Glows in dark	Does not glow in dark	
12. Action with NaOH	Evolves phosphine	No action	

#### Question 14.

Why does nitrogen show catenation properties less than phosphorus?

Answer:

The catenation properties depend upon the strength of the element-element bond. The N-N bond strength is much weaker (due to the repulsion of lone pairs on nitrogen because of its small size) than the P-P bond strength, therefore, nitrogen shows catenation less than phosphorus.

#### Question 15.

Give one disproportionation reaction of phosphorus acid  $(H_3PO_3)$ . Answer:

Upon heating to about 573 K, phosphorus acid undergoes disproportionation as follows :

 $4 H_3 PO_3 \xrightarrow{573 \text{ K}} 3H_3 PO_4 + PH_3$ Phosphorus acid Phosphoric acid

Question 16.

Can  $\mathsf{PCI}_{\scriptscriptstyle 5}$  act as oxidising as well as a reducing agent? Justify.

Answer:

In general, the molecules of a substance can behave as a reducing agent if the central atom is in a position to increase its oxidation number. Similarly, they can act as an oxidising agent if the central atom is in a position to decrease its oxidation number. Now, the maximum oxidation state or oxidation number of phosphorus (P) is +5. It cannot increase the same but at the same time can decrease its oxidation number. In PCl<sub>5</sub>, oxidation number of P is already +5. It therefore, cannot act as a reducing agent. However, it can behave as an oxidising agent in certain reactions in which its oxidation number decreases. For example,

$$2Ag + \stackrel{+5}{PCl_5} \longrightarrow 2AgCl + \stackrel{+3}{PCl_3}$$
$$Sn + 2\stackrel{+5}{PCl_5} \longrightarrow SnCl_4 + 2\stackrel{+3}{PCl_3}$$

## Question 17. Justify the placement of O, S, Se, Te and Po in the same group of the periodic table in terms of

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electronic configuration, oxidation states and hydride formation. Answer:

The members of the oxygen family are placed in group 16 of p-block. Their inclusion in the same group is justified on the basis of the following characteristics.

1. Electronic configuration. Members of the family have ns2p4 configuration. Their group (10 + 6) is 16.

2. Oxidation states. With exception of oxygen which exhibits -2 oxidation state in its compounds  $(OF_2 \text{ and } H_2O_2 \text{ are exceptions})$ , rest of the members of the family show variable oxidation states (-2, +2, +4, +6) in their compounds.

3. Hydride formation. All the members of the family form covalent hydrides (MH<sub>2</sub>) in which the central atom is sp<sup>3</sup> hybridised. These have angular structures and their characteristics show regular gradation.

 $\begin{array}{c|ccccc} H_2O & H_2S & H_2Se & H_2Te & H_2Po \\ \hline Water & Hydrogen sulphide & Selenium hydride & Tellurium hydride & Polonium hydride \\ \hline FeS + H_2SO_4 (dil.) \rightarrow FeSO_4 + H_2S \\ Na_2Se + H_2SO_4 (dil.) \rightarrow Na_2SO_4 + H_2Se \end{array}$ 

Question 18.

Why is dioxygen a gas while sulphur is a solid? (C.B.S.E. Delhi 2013) Answer:

The oxygen atom has a tendency to form  $p\pi$ - $p\pi$  multiple bonding due to its small size and high electronegativity. As a result, oxygen exists as a diatomic molecule. These mol¬ecules are held together by weak van der Waal's forces of attraction which can be easily overcome by collisions of the molecules at room temperature. Therefore,  $O_2$  is the gas at room temperature.

Sulphur, on the other hand, because of its bigger size and lower electronegativity, does not form  $p\pi$ - $p\pi$  multiple bonds. Instead, it prefers to form an S-S single bond and form polyatomic complex molecules having eight atoms per molecule (S<sub>8</sub>) and have puckered ring structure. Therefore, S atoms are strongly held together and it exists as a solid.

Question 19.

Knowing the electron gain enthalpy values for  $0 \rightarrow 0^{-}$  and  $0 \rightarrow 0^{2}$  as – 141 kJ mol<sup>-1</sup> and 702 kJ mol<sup>-1</sup> respectively, how can you account for the formation of a large number of oxides having  $0^{2}$  species and not  $0^{-}$ ?

Answer:

The stability of an ionic compound depends on its lattice energy. The more the lattice energy of a compound, the 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 O<sup>2-</sup> ion is much more than the oxide involving O<sup>-</sup> ion. Hence, we can say that the formation of O<sup>2-</sup> is energetically more favourable than the formation of O<sup>-</sup>

Question 20.

Which aerosols deplete ozone?

Answer:

Aerosols or chlorofluorocarbons (CFC's) such as freon (CCl<sub>2</sub>F<sub>2</sub>) deplete ozone layer by supplying



$$CF_2Cl_2 \xrightarrow{hv} {}^{\bullet}CF_2Cl + Cl^{\bullet} \text{ (free radical)}$$

$$CFCl_3 \xrightarrow{hv} {}^{\bullet}CFCl_2 + Cl^{\bullet}$$

$$Cl^{\bullet} + O_3 \longrightarrow ClO^{\bullet} + O_2$$

$$ClO^{\bullet} \xrightarrow{slow} Cl^{\bullet} + {}^{\bullet}O$$

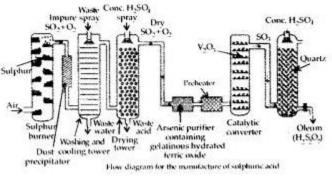
$$ClO^{\bullet} + {}^{\bullet}O \longrightarrow Cl^{\bullet} + O_2$$

Question 21. Describe the manufacture of  $H_2SO_4$  by Contact process. Answer:

Sulphuric acid is manufactured by the Contact process which involves three steps :

- 1. burning of sulphur or sulphide ores in air to generate SO<sub>2</sub>.
- 2. conversion of SO<sub>2</sub> to SO<sub>3</sub> by the reaction with oxygen in the presr nee of a catalyst  $(V_2O_5)$ , and
- 3. absorption of SO<sub>3</sub> in  $H_2SO_4$  to give oleum ( $H_2S_2O_7$ ).

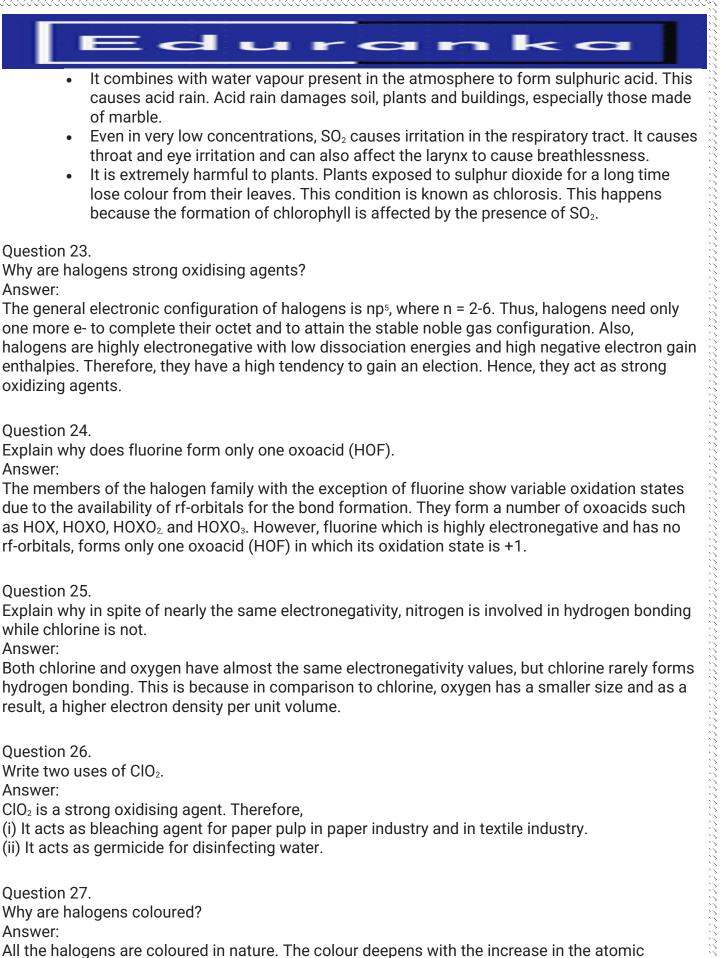
A flow diagram for the manufacture of sulphuric acid is shown in the figure. The SO<sub>2</sub> produced is purified by removing dust and other impurities such as arsenic compounds. The key step in the manufacture of  $H_2SO_4$  is the catalytic oxidation of SO<sub>2</sub> with O<sub>2</sub> to give SO<sub>3</sub> in the presence of  $V_2O_5$  (catalyst).



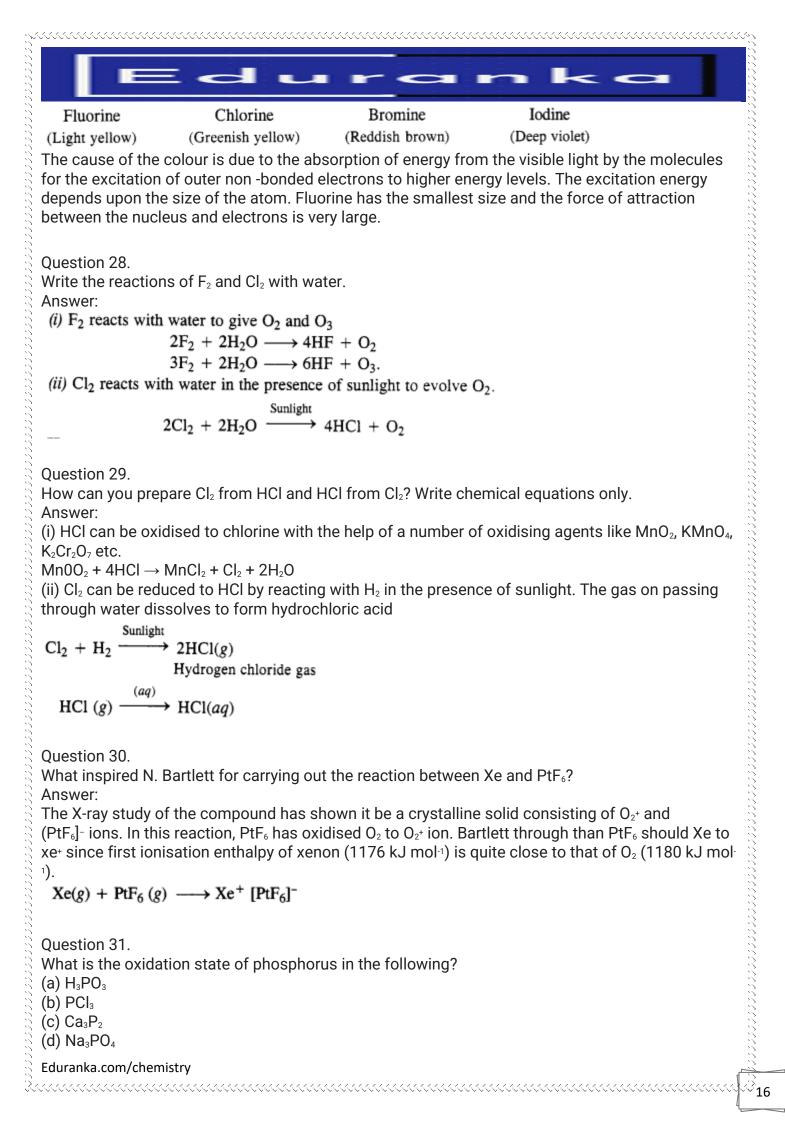
The reaction is exothermic, reversible and the forward reaction leads to a decrease in volume. Therefore, low temperature and high pressure are favourable conditions for maximum yield. But the temperature should not be very low otherwise rate of reaction will become slow.

In practice, the plant is operated at a pressure of 2 bar and a temperature of 720 K. The SO<sub>3</sub> gas from the catalytic converter is absorbed in concentrated  $H_2SO_4$  to produce oleum. Dilution of oleum with water gives  $H_2SO_4$  of the desired concentration. In the industry, two steps are carried out simultaneously to make the process a continuous one and also to reduce the cost.

Question 22. How is  $SO_2$  an air pollutant? Answer:



number of the element from fluorine to iodine.



#### (e) POF<sub>3</sub> Answer:

 $H_3PO_3: H_3 \stackrel{+1}{P} \stackrel{x}{O_3} \stackrel{-2}{O_3}$ (a) 3(+1) + x + 3(-2) = 0 or x - 3 = 0or + 3 (b)  $PCl_3 : \stackrel{x}{P} \stackrel{-1}{Cl_3}$ x + 3(-1) = 0 or x - 3 = 0or + 3 (c)  $Ca_3P_2: \overset{+2}{C}a_3\overset{x}{P}_2$ 3(+2) + 2(x) = 0 or 2x + 6 = 0or - 3 (d)  $Na_3PO_4: Na_3PO_4^{+1} PO_4^{x-2}$ 3(+1) + x + 4(-2) = 0 or x - 5 = 0or + 5 (e) **POF<sub>3</sub>** :  $\overset{x}{P} \overset{-2}{O} \overset{-1}{F_3}$ x + (-2) + 3(-1) = 0or x - 5 = 0or + 5

Question 32.

Write balanced equations for the following :

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

(i) NaCl + H<sub>2</sub>SO<sub>4</sub>  $\longrightarrow$  NaHSO<sub>4</sub> + HCl] × 4  $\frac{4HCl + MnO_2 \longrightarrow MnCl_2 + Cl_2 + 2H_2O}{4NaCl + MnO_2 + 4H_2SO_4 \longrightarrow 4NaHSO_4 + MnCl_2 + Cl_2 + 2H_2O}$ (ii) Cl<sub>2</sub>(g) + 2NaI(aq)  $\longrightarrow 2NaCl(aq) + I_2(s)$ 

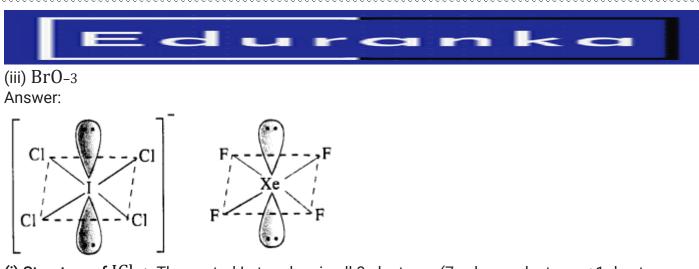
Question 33.

How are xenon fluorides  $XeF_2$ ,  $XeF_4$  and  $XeF_6$  obtained ? Answer:

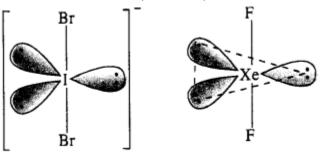
Xenon forms three binary fluorides,  $XeF_2$ ,  $XeF_4$  and  $XeF_6$  by the direct reaction of elements under appropriate experimental conditions.

Nickel tube  $Xe(g) + F_2(g)$  $XeF_2(s)$ 675 K, 1 bar (2 vol) (1 vol) Xenon difluoride Nickel tube  $Xe(g) + 2F_2(g)$  $XeF_4(s)$ 873K, 7bar (1 vol) (5 vol) Xenon tetrafluoride Nickel tube  $Xe(g) + 3F_2(g)$  $\rightarrow$  XeF<sub>6</sub>(s) 575K, 60-70 bar (1 vol) (20 vol) Xenon hexafluoride

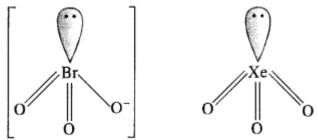
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Answer: $OF_2$ and CIF are isoelectronic to CIO-, out of which CIF is a Lewis base.
Question 35. How are XeO <sub>3</sub> and XeOF <sub>4</sub> prepared? Answer: Preparation of XeO <sub>3</sub> . By complete hydrolysis of XeF <sub>6</sub> . XeF <sub>6</sub> + $3H_2O \rightarrow XeO_3$ + 6HF. Preparation of XeOF <sub>4</sub> . By partial hydrolysis of XeF <sub>6</sub> . XeF <sub>6</sub> + $H_2O \rightarrow XeOF_4$ + 2HF.
Question 36 Arrange the following in the order of property indicated for each set: (i) $F_2$ , $Cl_2$ , $Br_2$ , $l_2$ – increasing bond dissociation enthalpy. Answer: Bond dissociation energy usually decreases on moving down a group as the atomic size increases. However, the bond dissociation energy of $F_2$ is lower than that of $Cl_2$ and $Br_2$ . This is due to the small atomic size of fluorine, Thus, the increasing order for bond dissociation energy among halogens is as follows: $l_2 < F_2 < Br_2 < Cl_2$
(ii) HF, HCl, HBr, HI – increasing acid strength. Answer: HF < HCl < HBr < HI The bond dissociation energy of HX molecules where X – F, Cl, Br, I, decreases with an increase in the atomic size. Since H – I bond is the weakest, HI is the strongest acid.
(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: BiH <sub>3</sub> < SbH <sub>3</sub> < AsH <sub>3</sub> < PH <sub>3</sub> < NH <sub>3</sub> 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.
increases while the electron density on the atom decreases. Thus, the basic strength decreases. Question 37. Which one of the following does not exist? (i) XeOF <sub>4</sub> (ii) NeF <sub>2</sub> (iii) XeF <sub>6</sub> . Answer: NeF <sub>2</sub> does not exist because the element Ne(Z = 10) with 1s <sup>2</sup> 2s <sup>2</sup> 2p <sup>6</sup> does not have vacant 2d orbitals. As such, there is no scope of any electron promotion even by a highly electronegative element. Question 38. Give the formula and describe the structure of a noble gas species which is isostructural with : (i) ICl-4
Question 38. Give the formula and describe the structure of a noble gas species which is isostructural with : (i) ICl-4 (ii) IBr-2 Eduranka.com/chemistry



(i) Structure of ICl-4. The central I atom has in all 8 electrons (7 valence electrons +1 due to negative charge). Out of these, it shares 4 electrons with four atoms of Cl and the remaining four electrons constitute two lone pairs. In all, there are six pairs. The structure of the ion must be octahedral or distorted square planar in order to minimise the forces of repulsion among the two electrons pairs. ICl-4 has  $(7 + 4 \times 7 + 1) = 36$  valence electrons and it iso-electronic and iso-structural with XeF<sub>4</sub> (8 + 4 x 7) which has also 36 valence electrons.



(ii) Structure of IBr-2. In IBr-2 ion, the central I atom has 8 valence electrons (7 + 1). Out of these, it shares 2 electrons with two atoms of Br and the remaining 6 electrons constitute three lone pairs. In all, there are five pairs. The structure of the ion must be trigonal bipyramidal or linear in order to minimise the force of repulsion among the three lone electrons pairs. IBrf has (7 + 2 x 7 + 1) = 22 valence electrons and is isoelectronic as well as iso-structural with noble gas species XeF<sub>2</sub> which has also 22 (8 + 2 x 7) electrons.



(iii) Structure of  $BrO_{-3}$  ion. In  $BrO_{-3}$  ion, the central Br atom has 8 valence electrons (7 + 1). Out of these, it shares 4 with two atoms of 0 forming Br=0 bonds. Out of the remaining four electrons, 2 are donated to the third 0 atom and account for its negative charge. The remaining 2 electrons constitute one lone pair. In order to minimise the force of repulsion, the structure of  $BrO_{-3}$  ion must be pyramidal.  $BrO_{-3}$  ion has (7 + 3 x 6 + 1) = 26 valence electrons and is isoelectronic as well as iso-structural with noble gas species Xe03 which has also 26 (8 + 3 x 6) electrons. Question 39.

Why do noble gases have comparatively large atomic sizes? Answer:

Noble gases do not form molecules. In the case of noble gases. the atomic radii correspond to

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Van der Waa1s radii. On the other hand, the atomic radii of other elements correspond to their covalent radii. By definition, Yan 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.

Question 40. List the uses of neon and argon gases. Answer: Uses of Neon:

- Neon lights are used for commercial advertisements.
- It consists of a long tube fitted with electrodes at both ends.
- On filling the tube with neon gas and passing electric discharge of about 1000 volt potential, a bright red light is produced.
- Different colours can be obtained by mixing neon with other gases.
- For producing blue or green light neon is mixed with mercury vapours.

Uses of Argon:

- Like helium, it is also used to create an inert atmosphere in welding aluminium and stainless steel.
- It is filled in electric bulbs along with 25% nitrogen.
- It is also used in radio valves.
- Argon alone or its mixture with neon is used in tubes for producing lights of different colours.

We hope the NCERT Solutions for Class 12 Chemistry Chapter 7 The p-Block Elements help you. If you have any query regarding NCERT Solutions for Class 12 Chemistry Chapter 7 The p-Block Elements, drop a comment below and we will get back to you at the earliest.