

# NCERT SOLUTIONS

## CLASS-XII CHEMISTRY

### CHAPTER-7

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## THE P BLOCK ELEMENTS

**Q 1: Briefly describe the general characteristics of Group 15 elements with reference to their oxidation state, electronic configuration, atomic size, electronegativity and ionisation enthalpy.**

**Answer**

**General trends in group 15 elements**

**(i) Electronic configuration:** There are 5 valence electrons for all the elements in group 15.

$ns^2np^3$  is their general electronic configuration.

**(ii) Oxidation states:** All these elements require three or more electrons to complete their octets and have 5 valence electrons. It is difficult in gaining electrons as the nucleus will have to attract three more electrons. This happens 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**

Ionization decreases as we move down the group. This happens because of increase in atomic sizes. Moving down the group, electronegativity decreases due to increase in size.

**(iv) Atomic size:** As we move down the group atomic size increases. This increase in the atomic size is attributed to an increase in the number of shells.

**Q 2: 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 unable to form  $p\pi - p\pi$  bonds with itself. This property is not exhibited by atoms such as phosphorus. Thus, phosphorus is more reactive than nitrogen.

**Q 3: 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.

**Q 4: Why does  $NH_3$  form hydrogen bond but  $PH_3$  does not?**

**Answer**

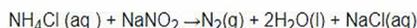
When compared to phosphorus nitrogen is highly electronegative. This results in a greater

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  $NH_3$ .

**Q 5: 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.

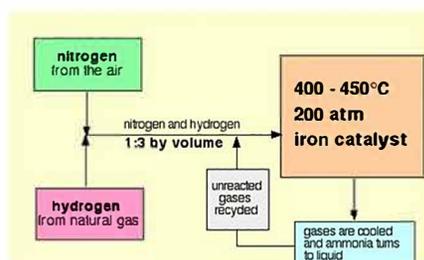
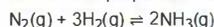


NO and  $HNO_3$  are produced in small amounts. These are impurities that can be removed on passing nitrogen gas through aqueous sulphuric acid, containing potassium dichromate.

**Q 6: How is ammonia manufactured industrially?**

**Answer**

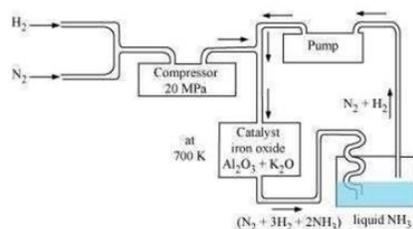
Ammonia is prepared on a large-scale by the Haber's process.



liquid ammonia

The optimum conditions for manufacturing ammonia are:

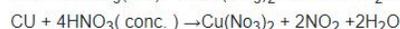
- (i) Pressure (around  $200 \times 10^5$  Pa)
- (ii) Temperature (4700 K)
- (iii) Catalyst such as iron oxide with small amounts of  $Al_2O_3$  and  $K_2O$



**Q 7: Illustrate how copper metal can give different products on reaction with  $HNO_3$ .**

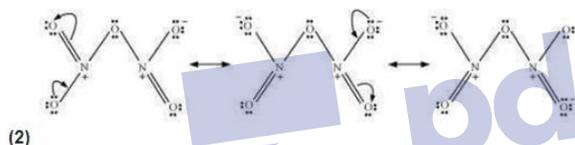
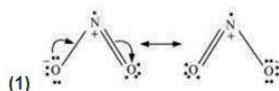
**Answer**

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



**Q 8: Give the resonating structures of  $NO_2$  and  $N_2O_5$ .**

**Answer**



**Q 9: The HNH angle value is higher than HPH, HAsH and HSbH angles. Why? [Hint: Can be explained on the basis of  $sp^3$  hybridisation in  $NH_3$  and only s-p bonding between hydrogen and other elements of the group].**

**Answer**

Hydride  $NH_3$   $PH_3$   $AsH_3$   $SbH_3$

H-M-H angle  $107^\circ$   $92^\circ$   $91^\circ$   $90^\circ$

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.

**Q 10: Why does  $R_3P = O$  exist but  $R_3N = O$  does not ( $R = \text{alkyl group}$ )?**

**Answer**

N (unlike P) lacks the d-orbital. This restricts nitrogen to expand its coordination number beyond four. Hence,  $R_3N = O$  does not exist.

**Q 11: Explain why  $NH_3$  is basic while  $BiH_3$  is only feebly basic.**

**Answer**

**$NH_3$  is distinctly basic while  $BiH_3$  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.

**Q 12: Nitrogen exists as diatomic molecule and phosphorus as  $P_4$ . Why?**

**Answer**

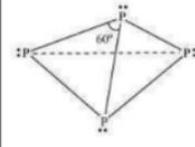
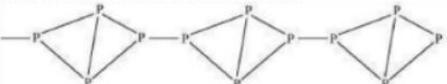
Nitrogen owing to its small size has a tendency to form  $p\pi-p\pi$  multiple bonds with itself.

Nitrogen thus forms a very stable diatomic molecule,  $N_2$ . On moving down a group, the tendency to form  $p\pi-p\pi$  bonds decreases (because of the large size of heavier elements). Therefore, phosphorus (like other heavier metals) exists in the  $P_4$  state.

**Q 13: Write main differences between the properties of white phosphorus and red phosphorus.**

**Answer :**

White phosphorus	Red phosphorus
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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 $P_4$ molecule. 	It exists as a chain of tetrahedral $P_4$ units. 

**Q 14: Why does nitrogen show catenation properties less than phosphorus?**

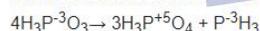
**Answer**

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.

**Q 15: Give the disproportionation reaction of  $H_3PO_3$ .**

**Answer**

On heating, orthophosphorous 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.



**Q 16: Can  $PCl_5$  act as an oxidising as well as a reducing agent? Justify.**

**Answer**

$PCl_5$  can only act as an oxidizing agent. The highest oxidation state that P can show is +5. In  $PCl_5$ , phosphorus is in its highest oxidation state (+5). However, it can decrease its oxidation state and act as an oxidizing agent.

**Q 17: 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 n varies from 2 to 6

(ii) **Oxidation state:**

As these elements have six valence electrons ( $ns^2np^4$ ), 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.

**Q 18: Why is dioxygen a gas but sulphur a solid?**

**Answer**

Oxygen is smaller in size when compared to sulphur. Since its size is small, it can form  $p\pi-p\pi$  bonds and form  $O_2(O \equiv O)$  molecule. Also, the intermolecular forces in oxygen are weak van der Waals', which cause it to exist as gas. On the other hand, sulphur does not form  $M_2$  molecule but exists as a puckered structure held together by strong covalent bonds. Hence, it is a solid.

**Q 19: Knowing the electron gain enthalpy values for  $O \rightarrow O^{-1}$  and  $O \rightarrow O^{2-}$  as  $-141$  and  $702 \text{ kJ mol}^{-1}$  respectively, how can you account for the formation of a large number of oxides having  $O^{2-}$  species and not  $O^{-}$ ?**

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

**Answer**

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

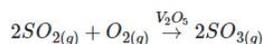
Lattice energy is directly proportional to the charge carried by an ion. When a metal combines with oxygen, the lattice energy of the oxide involving  $O^{2-}$  ion is much more than the oxide involving  $O^{-}$  ion. Hence, the oxide having  $O^{2-}$  ions are more stable than oxides having  $O^{-}$ . Hence, we can say that formation of  $O^{2-}$  is energetically more favourable than formation of  $O^{-}$ .

**Q 20. Which of the aerosol is responsible for the depletion of ozone?**

**Ans:**

The aerosol which is responsible for the depletion of ozone is: Freons or chlorofluorocarbons (CFCs)

The molecules of CFS breaks down when there is presence of ultraviolet radiations and forms chlorine free radicals which then combines with ozone to form oxygen.



**Q 21. Describe the manufacture of  $H_2SO_4$  by contact process?**

**Ans:**

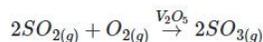
The steps which are required in the production of Sulphuric Acid by the contact process

Step (1)

Sulphide ores or Sulphur are burnt in air to form  $SO_2$ .

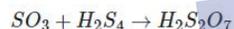
Step (2)

By a reaction with oxygen,  $SO_2$  is converted into  $SO_3$  in the presence of  $V_2O_5$  as a catalyst.



Step (3)

$SO_3$  produced is absorbed on  $H_2SO_4$  to give  $H_2S_2O_7$  (oleum).



This oleum is then diluted to obtain  $H_2SO_4$  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.

**Q 22: How does  $SO_2$  cause air pollution?**

**Soln:** The environment is harmed by sulphur dioxide in many ways:

1. Sulphuric acid is formed, when it is combined with water vapour present in the atmosphere. This causes acid that damages plants, soil, buildings (those made of marble are more prone) etc.
2.  $SO_2$  causes irritation in respiratory tract, throat, eyes and can also affect the larynx to cause breathlessness.
3. The color of the leaves of the plant gets faded when it is exposed to sulphur dioxide for a long time. This defect is known as chlorosis. The formation of chlorophyll is affected by the presence of sulphur dioxide.

**Q 23: Halogens are strong oxidizing agents. Explain.**

**Soln:** Halogens have an electronic configuration of  $np^5$ , where  $n=2-6$ . Thus, halogens require only one more electron to complete their octet and to attain the stable noble gas configuration. Moreover, halogens have high negative electron gain enthalpies and are highly electronegative with low dissociation energies. As a result, they have a high tendency to gain an electron. Hence, they act as strong oxidising agents.

**Q 24: Fluorine forms only one oxoacid, HOF. Explain.**

**Soln:** Fluorine has high electronegativity and a small size, hence it forms only one oxoacid i.e HOF.

**Q 25: Oxygen and Chlorine has nearly the same electronegativity, still oxygen forms hydrogen bonds while chlorine does not. Why?**

**Soln:** Oxygen has a smaller size and due to which a higher electron density per unit volume. Hence, oxygen forms hydrogen bonds while chlorine does not despite having the similar electronegative values.

**Q 26. State two applications of  $ClO_2$**

**Ans:** Applications of  $ClO_2$

- ( a )Used for purification of water.  
( b ) Used for bleaching.

**Q 27. What is the reason for halogens being colored?**

Ans : Halogens are colored because they take in radiations from the visible spectrum. This excites the valence electrons to a higher energy level. The amount of energy required for excitation differs from halogen to halogen, thus they exhibit different colors.

**Q28. Give the reactions of  $Cl_2$  and  $F_2$  with water.**

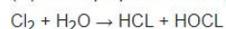
Ans: ( i )  $Cl_2 + H_2O \rightarrow HCl + HOCl$

( ii )  $2 F_2 + 2H_2O \rightarrow 4H^+ + 4F^- + O_2 + 4HF$

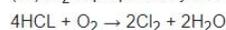
**Q29. Write the reactions involved in the preparations of HCl from  $Cl_2$  and  $Cl_2$  from HCl ?**

Ans:

( i ) HCl is prepared from  $Cl_2$  by reacting it with water.



( ii )  $Cl_2$  is prepared by Deacon's process from HCl



**Q30. What was the inspiration for N. Bartlett to carry out the reaction between  $PtF_6$  and Xe?**

Ans: N. Bartlett observed that  $PtF_6$  and  $O_2$  reacts to produce a compound  $O_2^+[PtF_6]^-$ .

As the first ionization enthalpy of Xe ( 1170 kJ/mol ) is very close to that of  $O_2$ , he figured that  $PtF_6$  could also oxidize Xe to  $Xe^+$ . Thus, he reacted  $PtF_6$  and Xe to form a red colored compound  $Xe^+[PtF_6]^-$ .

**Q31. In the compounds given below, find the oxidation states of phosphorus:**

( a )  $H_3PO_3$

( b )  $PCl_3$

( c )  $Ca_3P_2$

( d )  $Na_3PO_4$

( e )  $POF_3$ ?

Ans: Let the oxidation state of phosphorous be x

(a)  $H_3PO_3$

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

$$x - 3 = 0$$

$$x = 3$$

(b)  $PCl_3$

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

$$x = 3$$

(c)  $Ca_3P_2$

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

$$2x = -6$$

$$x = -3$$

(d)  $Na_3PO_4$

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

$$x - 5 = 0$$

$$x = 5$$

(e)  $POF_3$

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

$$x - 5 = 0$$

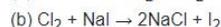
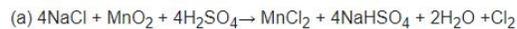
$$x = 5$$

**Q 32. Give balanced equations for the reactions below:**

( a ) NaCl being heated with  $H_2SO_4$  in the presence of  $MnO_2$ .

( b ) Chlorine gas passed through a Nalund water solution.

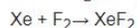
Ans:



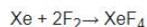
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**Q33. State the reaction that gives xenon fluorides  $\text{XeF}_2$ ,  $\text{XeF}_4$  and  $\text{XeF}_6$ .**

Ans:  $\text{XeF}_2$ ,  $\text{XeF}_4$  and  $\text{XeF}_6$  are obtained through direct reactions between Xe and  $\text{F}_2$ . The product depends upon the conditions of the reaction :



(excess)



( 1:5 ratio )



(1 : 20 ratio)

**Q34. Give the neutral molecule with which  $\text{ClO}^-$  isoelectronic. Is this molecule a Lewis base?**

Ans:  $\text{ClO}^-$  is isoelectronic with  $\text{ClF}$ .

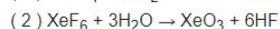
Total electrons in  $\text{ClO}^- = 17 + 8 + 1 = 26$

Total electrons in  $\text{ClF} = 17 + 9 = 26$

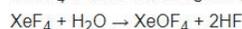
As  $\text{ClF}$  accepts electrons from F to form  $\text{ClF}_3$ ,  $\text{ClF}$  behaves like a Lewis base.

**Q35. State the preparation reaction of  $\text{XeO}_3$  and  $\text{XeOF}_4$ .**

Ans:  $\text{XeO}_3$  can be obtained using two methods :



$\text{XeOF}_4$  is obtained using  $\text{XeF}_6$



**Q36. Rearrange the given sets in the order as mentioned :**

( 1 )  $\text{Cl}_2$ ,  $\text{F}_2$ ,  $\text{I}_2$ ,  $\text{Br}_2$  – increasing bond dissociation enthalpy.

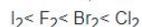
( 2 )  $\text{HCl}$ ,  $\text{HI}$ ,  $\text{HBr}$ ,  $\text{HF}$  – increasing acidic strength.

( 3 )  $\text{PH}_3$ ,  $\text{NH}_3$ ,  $\text{AsH}_3$ ,  $\text{BiH}_3$ ,  $\text{SbH}_3$  – increasing base strength.

Ans:

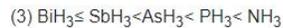
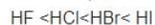
(1) Bond dissociation energy normally lowers on moving down a group because of increase in the atomic size. However,  $\text{F}_2$  has a lower bond dissociation energy than  $\text{Cl}_2$  and  $\text{Br}_2$ . This is because the atomic size of fluorine is very small.

Therefore, the increasing order for bond dissociation enthalpy is:



(2) Bond dissociation energy of a H-X molecule ( where X = F, Cl, Br, I ) lowers with an increase in the size of an atom. As, H-I bond is the weakest it will be the strongest acid.

Therefore, the increasing order acidic strength is :



On moving from nitrogen to bismuth, the atomic size increases but the electron density of the atom decreases. Hence, the basic strength lowers.

**Q37. Identify the one that does not exist, from among the following.**

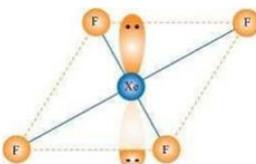


Ans: The one that does not exist is  $\text{NeF}_2$ .

**Q38. Present the structure and write the formula of a noble gas species that is isostructural with:**



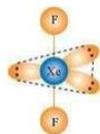
Ans:



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(a)  $\text{XeF}_4$  is isoelectronic to  $\text{Cl}_4^-$ . And it square planar in geometry :

(b)  $\text{XeF}_2$  is isoelectronic with  $\text{IBr}_2^-$ . It has a linear structure.



(c)  $\text{XeO}_3$  is isoelectronic and isostructural to  $\text{BrO}_3^-$ . It has a pyramidal structure.



**Q39. What is the reason for noble gases having relatively bigger atomic sizes ?**

Ans:

Noble gases have atomic radii that corresponds to van der Waal's radii. Whereas, other elements have a covalent radii. Now, by definition van der Waal's radii are bigger than covalent radii. This is the reason why noble gases have relatively bigger atomic sizes.

**Q40. Give some uses of argon and neon gases.**

Ans: Uses of Argon gas:

- (a) Argon is used to keep an inert atmosphere in high temperature metallurgical operations like arc welding.
- (b) It is used in fluorescent and incandescent lamps where it is required to check the sublimation of the filament. Thereby, increasing the life of the lamp.
- (c) Argon is used in laboratories to handle substances that are air-sensitive.

Uses of neon gas:

- (a) Neon is filled in discharge tubes for advertising or decoration.
- (b) Neon is used for making beacon lights.
- (c) It is used alongside helium to protect electrical equipments against high voltage.