

## CHAPTER - 15

# THE p-BLOCK ELEMENTS (Groups 13 - 18)

### SYNOPSIS

The p-block of the periodic table is unique in terms of having all types of elements-metals, non-metals and metalloids. There are six groups of p-block elements in the periodic table numbering from 13 to 18. Their valence shell electronic configuration is  $ns^2np^{1-6}$  except for He. Difference in the inner core of their E. C. greatly influence their physical & chemical properties. As a consequence of this, a lot of variation in properties among the elements is observed. In addition to the group oxidation state, these elements show other oxidation states differing from the total number of valence  $e^-$  by unit of two. While the group oxidation state is the most stable for the lighter elements of the group, the lower oxidation state becomes progressively more stable for the heavier elements. The combined effect of size and availability of d-orbitals considerably influences the ability of these elements to form  $\pi$ -bonds, while the lighter elements form  $p\pi - p\pi$  bonds, the heavier ones form  $d\pi - p\pi$  or  $d\pi - d\pi$  bonds. Absence of d-orbital in second period of elements limits their maximum covalence to four while heavier ones can exceed their limit.

#### **Elements of group-13 (B, Al, Ga, In, Tl)**

Boron is a typical non-metal and the other members are metals. The availability of three valence electrons for covalent bond formation using four orbitals ( $2s$ ,  $2p_x$ ,  $2p_y$  and  $2p_z$ ) leads to the so called electron deficiency in boron compounds. This deficiency makes them good electron acceptor and thus boron compounds behave as Lewis acids. Boron forms covalent molecular compounds with dihydrogen as boranes, the simplest of which is diborane ( $B_2H_6$ ). Diborane contains two bridging hydrogen atoms between two boron atoms. These bridge bonds are considered to be three-centre two electron bonds. The important compounds of boron are boric acid and borax. Boric acid,  $B(OH)_3$  is a weak monobasic acid; it acts as Lewis acid by accepting electrons from hydroxyl ions. Borax is a white crystalline solid of the formula  $Na_2[B_4O_5(OH)_4] \cdot 8H_2O$ . In borax bead test characteristic colours are produced by transition metals salts.

Aluminium exhibits +3 oxidation state. With heavier elements +1 oxidation state get progressively stabilised on going down the group. This is a consequence of inert pair effect.

#### **Elements of group-14 (C, Si, Ge, Sn, Pb)**

Carbon is a typical non-metal forming covalent bonds employing all its four valence electrons. It shows the property of catenation, the ability to form chains or rings, not only with C – C single bonds but also with multiple bonds ( $C = C$  or  $C \equiv C$ ). The tendency to catenation decreases as  $C \gg Si > Ge \approx Sn > Pb$ . Carbon provides one of the best examples of allotropy. Three important allotropes of carbon are diamond, graphite and fullerenes. The members of the carbon family mainly exhibit +4 and +2 oxidation states. Compounds in +4 oxidation state are generally covalent in nature. The tendency to show +2 oxidation state increases among heavier elements. Lead in +2 state is stable whereas in +4 oxidation state, it is a strong oxidising agent. Carbon also exhibits negative oxidation state. It forms two important oxides CO and  $CO_2$ . CO is neutral

whereas  $\text{CO}_2$  is acidic in nature. CO having lone pair of electrons on carbon forms metal carbonyls. It is deadly poisonous due to higher stability of its haemoglobin complex as compared to that of oxyhaemoglobin complex. Increased content of  $\text{CO}_2$  in the atmosphere is contributing to green house effect. Silica, silicates and silicones are important class of compounds and find applications in industry and Technology.

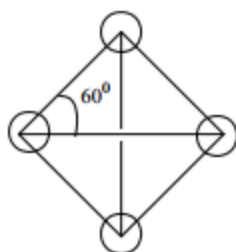
### Elements of Group 15 (N, P, As, Sb, Bi)

- $\text{N}_2$  is the major component (78% v/v) of atmospheric air, but in the mineral form it occurs as  $\text{NaNO}_3$  and  $\text{KNO}_3$  only, in the earth's crust. (Relative abundance is 33<sup>rd</sup>/78 for  $\text{N}_2$ ).
- Phosphorus occurs in the mineral apatite  $\text{Ca}_5(\text{PO}_4)_3\text{X}$  ( $\text{X} = \text{F}, \text{Cl}, \text{OH}$ ) and as phosphate in bones. (Relative abundance is 11<sup>th</sup>/78 for phosphorus)
- They have  $ns^2 sp^3$  configuration.
- They show +3, +5 and -3 oxidation states. Bi shows a stable positive oxidation state of +3 only (inert pair effect) N has oxidation states from -3 to +5. Bi shows -3 oxidation state in compounds with metals.
- They have exceptionally high  $\Delta_i H_1$  (compared with group 14 and group 16) except Bi
- Nitrogen is a diatomic gas, whereas other elements are tetraatomic solids.
- Reactivity :  $\text{N}_2$  is least reactive at room temperature due to a high bond-enthalpy of  $\text{N} \equiv \text{N}$  ( $941 \text{ kJ mol}^{-1}$ ) white phosphorus is the most reactive at room temperature due to a greater angular strain in  $\text{P}_4$  - tetrahedral structure.
- They form oxides of the formulae  $\text{E}_2\text{O}_3$  and  $\text{E}_2\text{O}_5$ . Their oxides show acidic character and acidity decreases from top to bottom in the group
- They form halides of the formulae  $\text{EX}_3$  and  $\text{EX}_5$ . Nitrogen does not form a pentahalide due to absence of d-orbitals. Among the halides of nitrogen, only  $\text{NF}_3$  is stable.
- They form binary compounds with metals, exhibiting -3 oxidation state. (eg:  $\text{Ca}_3\text{P}_2$ ,  $\text{Zn}_3\text{P}_2$ ,  $\text{Li}_3\text{N}$ ,  $\text{Mg}_3\text{N}_2$ ,  $\text{Na}_3\text{As}_2$ ,  $\text{Zn}_3\text{Sb}_2$ ,  $\text{Mg}_3\text{Bi}_2$  etc)
- They form hydrides of formula  $\text{EH}_3$ .
  - Thermal stability : Decreases from  $\text{NH}_3$  to  $\text{BiH}_3$
  - Reducing property : Increases from  $\text{NH}_3$  to  $\text{BiH}_3$
  - H - E - H bond angle : Decreases from  $\text{NH}_3$  to  $\text{BiH}_3$
  - Bond - enthalpy : Decreases from  $\text{NH}_3$  to  $\text{BiH}_3$
  - Boiling point :  $\text{PH}_3 < \text{AsH}_3 < \text{NH}_3 < \text{SbH}_3 < \text{BiH}_3$
  - Melting point :  $\text{PH}_3 < \text{AsH}_3 < \text{SbH}_3 < \text{NH}_3$
  - Lewis base character : Decreases from  $\text{NH}_3$  to  $\text{BiH}_3$
- $\text{NH}_3$  has a higher boiling point than  $\text{PH}_3$  and  $\text{AsH}_3$  due to strong hydrogen bonding.
- $\text{NH}_3$  has a larger bond angle due to high electronegativity of N. As electronegativity of central atom decreases bond angle decreases.

### VI. Phosphorus : Phosphorus is extracted from rock phosphate $\text{Ca}_3(\text{PO}_4)_2$

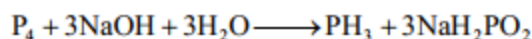
The important allotropes of phosphorus are white, red and black.

**White phosphorus :** It is a white waxy solid, insoluble in water and soluble in  $\text{CS}_2$ . It has phosphorescence in the dark. White phosphorus has tetrahedral molecular structure.

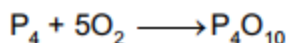


### Reaction with hot NaOH

It dissolves in boiling aq. NaOH in an inert atmosphere of  $\text{CO}_2$  forming phosphine ( $\text{PH}_3$ ) (preparation of  $\text{PH}_3$ )

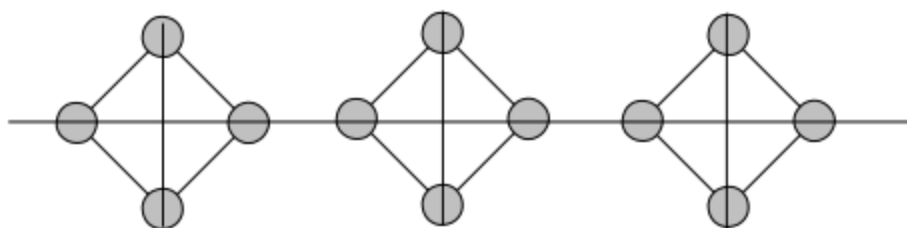


It burns in excess of air to form  $\text{P}_4\text{P}_{10}$



**Red phosphorus** : When white phosphorus is heated in an inert atmosphere at 573 K for seven days in a sealed cast-iron tube, red phosphorus is formed.

It is odourless, non poisonous, insoluble in  $\text{CS}_2$  or  $\text{H}_2\text{O}$ , less reactive, dense and polymeric.



**Black phosphorus** : It has two forms namely  $\alpha$  and  $\beta$ .  $\alpha$ -black phosphorus is formed when red P is heated in a sealed tube at 803 K. It is opaque having monoclinic or rhombohedral crystals.

It does not oxidise in air.  $\beta$ -black P is formed by heating white P at 473 K under high pressure. It does not burn in air when heated upto 673K.

### Elements of Group 16 (O, S, Se, Te, Po)

1. Occurrence : The most abundant element in the earth's crust is oxygen. The second most abundant element of group 16 is sulphur. Elements of group 16 are collectively called chalcogens since they are associated with ores (minerals). Se and Te form binary compounds with metals and they are found along with sulphides of copper and silver. Po occurs as a radioactive decay product.
2. Electronic configuration :  $ns^2 np^4$
3. Atomic and ionic radii : Increase from top to bottom in the group.
4. Ionisation enthalpy : Decreases from top to bottom in the group. I.E. of group 16 is less than expected.
5. Electron gain enthalpy : Oxygen has the lowest negative electron gain enthalpy in this group. Sulphur has the highest negative  $\Delta_{eg}H$ .
6. Electronegativity : Oxygen is the second most electronegative element in the periodic table. E.N. decreases from O to Po.
7. Physical state : O and S are non metals. Se and Te are metalloids. Po is a radioactive metal with a half life period of 13.8 days.  $\text{O}_2$  is a diatomic gas whereas other elements form octa atomic molecules.
8. Chemical properties : They show +2, +4 and +6 oxidation states in addition to -2. Oxygen does not show +4 and +6 oxidation states. Stability of +6 oxidation state decreases from S to Po.
9. Hydrides : All elements of group 16 form hydrides of formula  $\text{H}_2\text{E}$ . Stability of hydrides decreases from  $\text{H}_2\text{O}$  to  $\text{H}_2\text{Po}$ .  
 $\text{H}_2\text{S}$  has unexpectedly low MP and BP whereas  $\text{H}_2\text{O}$  has the highest MP and BP among the hydrides. Bond angle decreases from  $\text{H}_2\text{O}$  to  $\text{H}_2\text{Te}$ .
10. Oxides : They form oxides of formulae  $\text{EO}_2$  and  $\text{EO}_3$  except  $\text{O}_2$  (oxygen has no oxide).  $\text{SO}_2$  and  $\text{SO}_3$  are gases.  $\text{SeO}_2$  is a solid. Reducing property decreases from  $\text{SO}_2$  to  $\text{TeO}_2$ . ( $\text{SO}_2$  is a reducer whereas  $\text{TeO}_2$  is an oxidiser). Their trioxides are more acidic than dioxides.

11. Halides : They form halides of the type  $EX_2$ ,  $EX_4$  and  $EX_6$  except  $O_2$ .  
Among hexahalides, hexafluorides are stable. All hexafluorides are gases with octahedral structure.  
Among the tetrafluorides,  $SF_4$  is a gas,  $SeF_4$  is a liquid and  $TeF_4$  is a solid. They have see-saw geometry with  $sp^3d$  hybridisation of the central atom.  
They form dichlorides and dibromides, except oxygen. Their monohalides undergo dimerisation and tend to disproportionate (eg :  $2S_2Cl_2 \longrightarrow SCl_4 + 3S$ )

12. Oxides : Oxides are classified into simple oxides and mixed oxides. In simple oxides the element forming the oxide has only one oxidation state.

Eg :  $Na_2O$ ,  $MgO$ ,  $Al_2O_3$  etc

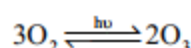
In mixed oxides the ox. states of the central atom are more than one.

Eg :  $Fe_3O_4 \rightleftharpoons FeO + Fe_2O_3$

Simple oxides are further classified into acidic, basic neutral and amphoteric oxides  $CO$ ,  $NO$  and  $N_2O$  are neutral oxides.

$P_4O_{10}$	$SO_3$	$Cl_2O_7$	etc	are acidic oxides
$ZnO$	$Al_2O_3$	$BeO$	etc	are amphoteric oxides
$Na_2O$	$CaO$	$BaO$	etc	basic oxides

13. Ozone ( $O_3$ ) : It is an allotrope of oxygen. In the upper atmosphere  $O_3$  is formed by the action of UV radiation.

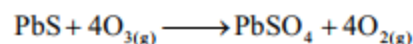


Preparation :  $O_3$  is prepared by subjecting pure dry  $O_2$  to a silent electric discharge.  $O_2$  containing about 10% (v/v)  $O_3$  is formed.



**Properties** :  $O_3$  is thermodynamically unstable. It is a pale-blue gas with a characteristic smell. It liquefies to give a dark blue liquid and solidifies to form violet-black solid.

It acts as a powerful oxidiser:



Nitric oxide reacts with ozone forming  $NO_2$  and  $O_2$



In this way  $NO$  causes depletion of  $O_3$

Ozone has an angular geometry with a bond angle of  $117^\circ$ .

Ozone is estimated by reacting with excess  $KI$  solution buffered with a borate and liberated  $I_2$  is quantitatively estimated using standard sodium thiosulphate solution.

**Uses** : Used as a germicide, disinfectant, for sterilising theatres and water, for bleaching oils, ivory, starch and in the oxidation of  $K_2MnO_4$ .

14. Sulphur : Sulphur has many isotopes, namely  $\alpha$ ,  $\beta$ ,  $\gamma$  etc.

$\alpha$ -sulphur or Rhombic sulphur : It is the stable allotrope of sulphur at room temperature. At 369 K it transforms into  $\beta$ -sulphur

$\beta$ -sulphur or monoclinic sulphur. It is stable above 369 K



Properties	$\alpha$ – sulphur	$\beta$ – sulphur
Molecular structure	$S_8$	$S_8$
M.P.	385.8 K	393 K
SP. gravity	2.06	1.98
Solubility in $CS_2$	Soluble	Soluble

Note : At about 1000 K, sulphur vapours contain  $S_2$  molecules. Which are paramagnetic like  $O_2$ .

### Elements of group 17 Halogens (F, Cl, Br, I and At)

- They form salts and so they are collectively called halogens (means salt producers)
- All halogens occur in the combined state and fluorine is the most abundant halogen in the earth's crust.  $Cl_2$  is the second most abundant halogen. Sea water contains chlorides, bromides and iodides.
- They have diatomic molecules.  $F_2$  and  $Cl_2$  are gases  $Br_2$  is a liquid and  $I_2$  is a sublimable solid at room temperature.
- They are strongly electronegative
- They have the shortest atomic radii in their respective periods due to high effective nuclear charge.
- They have the highest  $-\Delta_{eg}H_1$  in their respective periods. The highly  $-\Delta_{eg}H_1$  is due to attainment of stable electronic configuration after accepting one more electron by the atom. F has a lower  $-\Delta_{eg}H_1$  than Cl due to a higher electron-density of valence shell.
- Bond dissociation enthalpy is in the order :  $Cl-Cl > Br-Br > F-F > I-I$   
 Low bond dissociation enthalpy of  $F_2$  is due to strong  $\ell p - \ell p$  repulsion between the two F atoms bonded by a shorter bond. (143 pm)  
 Low bond-enthalpy, high electronegativity and higher hydration enthalpy of  $F^-$  make  $F_2$  more reactive among the non metals.
- Electronic configuration :  $ns^2 np^5$
- Oxidation state : +1, +3, +5, +7 and -1. Fluorine shows -1 oxidation state only at room temperature.

### Chemical properties

**(a) Reaction with Hydrogen** : They form hydrogen halides (HX) and reactivity towards hydrogen decreases from  $F_2$  to  $I_2$ .

**Variation of acidity** :  $HF < HCl < HBr < HI$  due to weakening of basic strength of halide ions in the order  $F^- > Cl^- > Br^- > I^-$  (when the conjugate base is strong the acid is weak.)

**Variation of b.p of HX** :  $HF > HI > HBr > HCl$ . HF has a high b.p. due to strong intermolecular hydrogen bonding and HI has the second highest b.p. due to strong vander waal's force

**Variation of M.P.** :  $HI > HF > HBr > HCl$

**(b) Reaction with  $O_2$**  : Most of the oxides of halogens are unstable.  $OF_2$  is a fluoride of  $O_2$  and act as a strong fluorinating agent. Another fluoride of  $O_2$ , namely  $O_2F_2$  is used to produce  $PuF_6$ .

$Cl_2$ ,  $Br_2$  and  $I_2$  form oxides in +1, +3, +5 and +7 oxidation states. The stability of oxides is in the order  $I > Cl > Br$  and oxides in higher oxidation state are more stable than in lower oxidation state.

Oxides of  $Cl_2$  ( $Cl_2O$ ,  $Cl_2O_6$  and  $Cl_2O_7$ ) are highly reactive oxidisers having explosive character.  $ClO_2$  is used to bleach paper - pulp and cotton and in water treatment.

Oxides of bromine are least stable and exist only at low temperatures. (middle row anomaly). (p-block element from Ga to Kr and d-block element from Y to Cd show anomalous properties due to a higher

effective nuclear charge resulting from the filling up of 3d and this is known as middle row-anomaly).

$I_2O_5$  is an oxidising agent and it is used in the estimation of CO.

**Reaction with metals** : They form halide with metals. Ionic character varies in the order.

$MF > MCl > MBr > MI$

Halides in higher O.S. are more covalent than the ones in lower O.S.

Eg :  $SnCl_4$ ,  $PbCl_4$ ,  $SbCl_5$  and  $UF_6$  are more covalent than  $SbCl_4$ ,  $PbCl_4$ ,  $SbCl_5$  and  $UF_6$  respectively

### Elements of group 18 (He, Ne, Ar, Kr, Xe and Rn)

**Occurrence** : All elements of group 18 except Rn occurs in the atmosphere. The most abundant noble gas in the universe is He and in the earth's atmosphere Ar is the most abundant noble gas. He is found associated with radioactive minerals and Rn is a decay product of Ra (226).

**Electronic configuration** :  $ns^2 sp^6$  except He

$\Delta_i H_1$  : Highest in the period

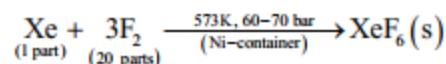
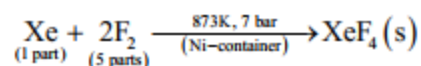
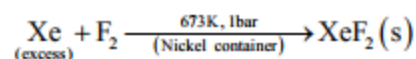
$\Delta_{eg} H$  : Highly positive

**Physical properties**: They are monoatomic gases. They are odourless, colourless and tasteless. They have very low MP and BP due to weak interatomic forces. Helium has the lowest B.P. among all known substances (4.2 K). Liquid helium can diffuse through glass rubber or plastics.

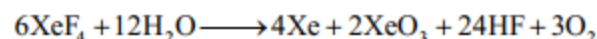
**Chemical properties** : They are chemically inert due to a *saturated valence shell electronic configuration*.

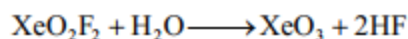
**Compounds of Noble-gases** : The first compound of any noble gas element prepared was  $XePtF_6$  (Neil Bartlett)

Xenon forms  $XeF_2$ ,  $XeF_4$  and  $XeF_6$  by direct combination with  $F_2$  under proper conditions.



They are colourless crystalline solids and sublime readily at 298 K. They are strong fluorinating agents. They undergo hydrolysis.

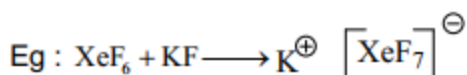
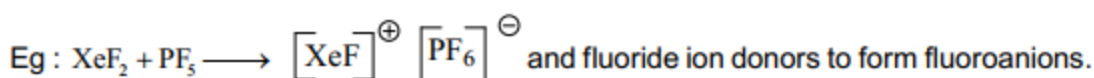




$\text{XeO}_3$  is a colourless explosive solid.

Compounds	Hybridisation state	Geometry
$\text{XeF}_2$	$\text{sp}^3\text{d}$	linear
$\text{XeF}_4$	$\text{sp}^3\text{d}^2$	square planar
$\text{XeF}_6$	$\text{sp}^3\text{d}^3$	distorted octahedral
$\text{XeO}_3$	$\text{sp}^3$	pyramidal
$\text{XeOF}_4$	$\text{sp}^3\text{d}^2$	square pyramidal

Xenon fluorides combine with fluoride ion acceptors to form cationic species.



**Uses :** Helium is used to inflate balloons, to produce very low temperature for cryogenic experiments, to dilute  $\text{O}_2$  for SCUBA to eliminate BENDS of deep sea divers. Also used to sustain powerful super conducting magnets used in MRI scanners.

Neon is used to produce red light in discharge tubes and sign-boards.

Argon is used to produce inert atmosphere in metallurgical processes and to fill incandescent bulbs. Xe and Kr are used in sign-boards.

### PART-I (JEE MAIN)

#### SECTION-I- Straight objective type questions

- Which of the following statements is false?
  - Boron has unusually high melting point due to very strong crystalline lattice
  - Gallium with unusually low melting point can exist as a liquid during summer
  - Radius of indium and thallium are comparable because thallium has noble gas plus 14 f-electrons plus 10 d-electron core
  - In group 13, electronegativity of the elements steadily decreases from boron to thallium

2. The stability of +1 oxidation state for group 13 elements increases in the sequence  
 1)  $Tl < In < Ga < Al$     2)  $In < Tl < Ga < Al$     3)  $Ga < In < Al < Tl$     4)  $Al < Ga < In < Tl$
3. The element that is used in high temperature thermometer is  
 1) Al                                  2) Hg                                  3) Ga                                  4) Pb
4. Which of the following lead halides does not exist?  
 1)  $PbF_4$                                   2)  $PbCl_4$                                   3)  $PbBr_4$                                   4)  $PbI_4$
5. Bond energy is highest for the bond  
 1)  $Sn-Sn$                                   2)  $C-C$                                   3)  $Si-Si$                                   4)  $Ge-Ge$
6. Which one of the following is not correct for the allotropic forms of carbon?  
 1)  $C_{60}$  molecule has a shape like soccer ball and its structure contains twenty six-membered rings and twelve five-membered rings  
 2) Graphite is thermodynamically the most stable allotrope of carbon  
 3) Graphite has a layered structure. Each layer is composed of hexagonal rings of carbon atoms  
 4) In diamond, each carbon atom is  $sp^2$  hybridised where as in graphite and fullerene, all the carbon atoms are  $sp^3$  hybridised
7. Which of the following statements is not correct for group 15 elements?  
 1) Except nitrogen all elements show allotropy  
 2) The N-N single bond is stronger than P-P single bond, thus catenation capacity is greater for nitrogen  
 3) Both black phosphorus and red phosphorus are less reactive than white phosphorus  
 4) Boiling point increases from top to bottom upto antimony but melting point increases up to arsenic and then decreases
8. Which of the following is an acidic oxide?  
 1)  $As_2O_3$                                   2)  $P_2O_3$                                   3)  $Bi_2O_3$                                   4)  $Sb_2O_3$
9. The correct order of ease of formation of hydrides is  
 1)  $NH_3 > PH_3 > AsH_3 > BiH_3$                                   2)  $BiH_3 > AsH_3 > PH_3 > NH_3$   
 3)  $NH_3 > AsH_3 > PH_3 > BiH_3$                                   4)  $BiH_3 > PH_3 > AsH_3 > NH_3$
10. Which of the following options correctly represent the variation in the indicated properties for the group 16 hydrides?  
 i)  $H_2O < H_2S < H_2Se < H_2Te$ : Acidic strength  
 ii)  $H_2O > H_2S > H_2Se > H_2Te$ : Reducing nature  
 iii)  $H_2O > H_2S > H_2Se > H_2Te$ : Thermal stability  
 iv)  $H_2Te > H_2O > H_2Se > H_2S$ : Boiling point  
 1) i, iii and iv                                  2) ii, and iii                                  3) i and iii                                  4) i, ii, iii and iv



11. Which of the following acids contain peroxy linkage?  
 I.  $\text{H}_2\text{S}_2\text{O}_7$                       II.  $\text{H}_2\text{S}_2\text{O}_8$                       III.  $\text{H}_2\text{SO}_5$                       IV.  $\text{H}_2\text{SO}_3$   
 1) I and II only                      2) II and III only                      3) I, II and III only                      4) II, III and IV only
12. The correct order of decreasing stability of hexafluorides of group 16 elements is :  
 1)  $\text{TeF}_6 > \text{SeF}_6 > \text{SF}_6$       2)  $\text{SF}_6 > \text{TeF}_6 > \text{SeF}_6$       3)  $\text{SF}_6 > \text{SeF}_6 > \text{TeF}_6$       4)  $\text{TeF}_6 > \text{SF}_6 > \text{SeF}_6$
13. Which of the following sets of compounds cannot be hydrolysed easily?  
 1)  $\text{SiCl}_4$ ,  $\text{NCl}_3$  and  $\text{TeF}_6$     2)  $\text{CCl}_4$ ,  $\text{NF}_3$  and  $\text{SF}_6$     3)  $\text{GeCl}_4$ ,  $\text{PCl}_3$  and  $\text{TeF}_6$     4)  $\text{SnCl}_4$ ,  $\text{PCl}_5$  and  $\text{TeF}_6$
14. Which of the following reactions is non spontaneous?  
 1)  $\text{Cl}_2 + \text{H}_2\text{O} \rightarrow \text{HCl} + \text{HOCl}$                       2)  $2\text{I}_2 + 2\text{H}_2\text{O} \rightarrow 4\text{HI} + \text{O}_2$   
 3)  $\text{Br}_2 + \text{H}_2\text{O} \rightarrow \text{HBr} + \text{HOBr}$                       4)  $2\text{F}_2 + 2\text{H}_2\text{O} \rightarrow 4\text{HF} + \text{O}_2$
15. The high oxidising power of  $\text{F}_2$  is due to its  
 1) high electron affinity  
 2) low enthalpy of hydration and high enthalpy of dissociation  
 3) high enthalpy of hydration and low enthalpy of dissociation  
 4) high enthalpy of hydration and high enthalpy of dissociation
16. The correct order of boiling point of hydrogen halides is  
 1)  $\text{HF} > \text{HI} > \text{HBr} > \text{HCl}$       2)  $\text{HF} > \text{HCl} > \text{HI} > \text{HBr}$       3)  $\text{HF} > \text{HCl} > \text{HBr} > \text{HI}$       4)  $\text{HI} > \text{HF} > \text{HBr} > \text{HCl}$
17. Match the following :
- | Column-I   | Column-II                       |
|--|---------------------------------|
| A) $\text{I}_2 < \text{Br}_2 < \text{Cl}_2 < \text{F}_2$ | P) pKa                          |
| B) $\text{Cl}_2 > \text{Br}_2 > \text{F}_2 > \text{I}_2$ | Q) Standard reduction potential |
| C) $\text{HI} < \text{HBr} < \text{HCl} < \text{HF}$     | R) Bond energy                  |
| D) $\text{HI} > \text{HF} > \text{HBr} > \text{HCl}$     | S) Melting point                |
| 1) A-R, B-Q, C -P, D- S                                  | 2) A-Q, B-R, C - P, D - S       |
| 3) A-Q, B-R, C-S, D - P                                  | 4) A -R, B-Q, C-S, D - P        |
18. Which of the following is not true about helium?  
 1) It has the lowest boiling point of any known substance  
 2) It has the highest first ionization energy  
 3) It can diffuse through rubber, glass and plastic material  
 4) It is the least abundant element of the group

19. Match List I with List II and select the correct answer using the codes given below the list

**List-I (Compound)**

A.  $\text{XeF}_4$

B.  $\text{XeF}_6$

C.  $\text{XeO}_3$

D.  $\text{XeOF}_4$

**List-II (Molecular shape)**

1. Distorted octahedral

2. Square pyramidal

3. Square planar

4. Pyramidal

1)  $A \rightarrow 1$ ;  $B \rightarrow 2$ ;  $C \rightarrow 3$ ;  $D \rightarrow 4$

2)  $A \rightarrow 3$ ;  $B \rightarrow 1$ ;  $C \rightarrow 4$ ;  $D \rightarrow 2$

3)  $A \rightarrow 1$ ;  $B \rightarrow 3$ ;  $C \rightarrow 2$ ;  $D \rightarrow 4$

4)  $A \rightarrow 2$ ;  $B \rightarrow 4$ ;  $C \rightarrow 1$ ;  $D \rightarrow 3$

20. If  $\text{XeF}_2$  and  $\text{XeF}_6$  are hydrolysed separately, then

1) Both give out  $\text{O}_2$

2)  $\text{XeF}_6$  gives  $\text{O}_2$  but  $\text{XeF}_2$  does not

3)  $\text{XeF}_2$  alone gives out  $\text{O}_2$

4) Neither of them gives HF

**SECTION-II - Numerical Type Questions**

21. The atomic number of an element with highest first ionisation enthalpy in group 13 is .....

22. Lead is a strong oxidising agent in the oxidation state of .....

23. The P-P-P bond angle in white phosphorous is close to .....°

24. The total number of P-H bond(s) in  $\text{H}_3\text{PO}_2$  is .....

25. How many of the following do(es) not exist?

$[\text{SiF}_6]^{2-}$ ,  $[\text{GeCl}_6]^{2-}$ ,  $[\text{SiCl}_6]^{2-}$ ,  $[\text{BF}_6]^{3-}$ ,  $[\text{Al}(\text{H}_2\text{O})_6]^{3+}$ ,  $[\text{Al}(\text{OH})_4]^-$