CHAPTER

# The p-Block Elements

# 7.1 Group 15 Elements

- 1. In which of the following compounds, nitrogen exhibits highest oxidation state?
  - (a)  $N_2H_4$  (b)  $NH_3$
  - (c)  $N_3H$  (d)  $NH_2OH$  (2012)
- 2. Nitrogen forms  $N_2$ , but phosphorus does not form  $P_2$ , however, it converts  $P_4$ , reason is
  - (a) triple bond present between phosphorus atom
  - (b)  $p\pi p\pi$  bonding is weak
  - (c)  $p\pi p\pi$  bonding is strong
  - (d) multiple bonds form easily. (2001)

# Which of the following oxides is most acidic? (a) As<sub>2</sub>O<sub>5</sub> (b) P<sub>2</sub>O<sub>5</sub> (c) N<sub>2</sub>O<sub>5</sub> (d) Sb<sub>2</sub>O<sub>5</sub> (1999)

4. Which of the following has the highest dipole moment?(b) A - H

(a) $SbH_3$	(b) $AsH_3$	
(c) $NH_3$	(d) PH <sub>3</sub>	(1997)

- 5. The basic character of hydrides of the V group elements decreases in the order
  (a) NH<sub>3</sub> > PH<sub>3</sub> > AsH<sub>3</sub> > SbH<sub>3</sub>
  - (b)  $SbH_3 \ge AsH_3 > PH_3 \ge NH_3$ (c)  $SbH_3 \ge PH_3 \ge AsH_3 \ge NH_3$

(d) 
$$NH_3 > SbH_3 > PH_3 > AsH_3$$
 (1996)

6. Among the following oxides, the lowest acidic is

(a) As<sub>4</sub>O<sub>6</sub>
(b) As<sub>4</sub>O<sub>10</sub>
(c) P<sub>4</sub>O<sub>6</sub>
(d) P<sub>4</sub>O<sub>10</sub>
(1996)

7. Which of the following fluorides does not exist?

(a) NF <sub>5</sub>	(b) PF <sub>5</sub>	
(c) $AsF_5$	(d) $SbF_5$	(1993)

8. Which one has the lowest boiling point?
(a) NH<sub>3</sub>
(b) PH<sub>3</sub>
(c) AsH<sub>3</sub>
(d) SbH<sub>3</sub>
(1989)

#### 7.2 Dinitrogen

9. Number of electrons shared in the formation of nitrogen molecule is

(a) 6 (b) 10	
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(c) 2 (d) 8 (1992)

- **10.** Nitrogen is relatively inactive element because
  - (a) its atom has a stable electronic configuration
  - (b) it has low atomic radius
  - (c) its electronegativity is fairly high
  - (d) dissociation energy of its molecule is fairly high.

(1992)

- **11.** Pure nitrogen is prepared in the laboratory by heating a mixture of
  - (a)  $NH_4OH + NaCl$  (b)  $NH_4NO_3 + NaCl$
  - (c)  $NH_4Cl + NaOH$  (d)  $NH_4Cl + NaNO_2.(1991)$
- **12.** Which of the following statement is not correct for nitrogen?
  - (a) Its electronegativity is very high.
  - (b) *d*-orbitals are available for bonding.
  - (c) It is a typical non-metal.
  - (d) Its molecular size is small. (1990)

#### 7.3 Ammonia

- **13.** Urea reacts with water to form *A* which will decompose to form *B*. *B* when passed through  $Cu^{2+}_{(aq)}$ , deep blue colour solution *C* is formed. What is the formula of *C* from the following?
  - (a)  $CuSO_4$  (b)  $[Cu(NH_3)_4]^{2+}$ (c)  $Cu(OH)_2$  (d)  $CuCO_3 \cdot Cu(OH)_2$ (*NEET 2020*)
- **14.** Aqueous solution of ammonia consists of

(a) $H_{H^+}^+$	(a) $NH^{-}$ and $OH^{-}$ . (1991)
4	4

### 7.4 Oxides of Nitrogen

- 15. Which of the following oxides of nitrogen is paramagnetic?
  (a) NO<sub>2</sub>
  (b) N<sub>2</sub>O<sub>3</sub>
  (c) N<sub>2</sub>O
  (d) N<sub>2</sub>O<sub>5</sub>
  (1994)
  16. Which of the following is a nitric acid anhydride?
  (a) NO
  (b) NO<sub>2</sub>
  - (a)  $NO_2$ (c)  $N_2O_5$  (d)  $N_2O_3$  (1988)

### 7.5 Nitric Acid

**17.** When copper is heated with conc. HNO<sub>3</sub> it produces

- (a)  $Cu(NO_3)_2$ , NO and  $NO_2$
- (b)  $Cu(NO_3)_2$  and  $N_2O$  (c)  $Cu(NO_3)_2$  and  $NO_2$
- (d)  $Cu(NO_3)_2$  and NO (NEET-I 2016)
- **18.** Zn gives H<sub>2</sub> gas with H<sub>2</sub>SO<sub>4</sub> and HCl but not with HNO<sub>3</sub> because
  - (a) Zn act as oxidising agent when react with  $HNO_3$
  - (b)  $HNO_3$  is weaker acid than  $H_2SO_4$  and HCl
  - (c) in electrochemical series Zn is above hydrogen
  - (d) NO<sub>3</sub><sup>-</sup> is reduced in preference to hydronium ion. (2002)
- **19.** Sugarcane on reaction with nitric acid gives
  - (a)  $CO_2$  and  $SO_2$  (b)  $(COOH)_2$
  - (c) 2HCOOH(two moles)(d) no reaction. (1992)

#### 7.6 Phosphorus - Allotropic Forms

- **20.** Which of the following phosphorus is the most reactive?
  - (a) Scarlet phosphorus (b) White phosphorus
  - (c) Red phosphorus (d) Violet phosphorus

(1999)

- **21.** Each of the following is true for white and red phosphorus except that they
  - (a) are both soluble in  $CS_2$
  - (b) can be oxidised by heating in air
  - (c) consist of the same kind of atoms
  - (d) can be converted into one another. (1989)

# 7.7 Phosphine

- 22. A compound 'X' upon reaction with  $H_2O$  produces a colourless gas 'Y' with rotten fish smell. Gas 'Y' is absorbed in a solution of CuSO<sub>4</sub> to give Cu<sub>3</sub>P<sub>2</sub> as one of the products. Predict the compound 'X'.
  - (a)  $Ca_3P_2$ (b)  $NH_4Cl$ (c)  $As_2O_3$ (d)  $Ca_3(PO_4)_2$ (Odisha NEET2019)
- **23.**  $PH_4I + NaOH$  forms (a)  $PH_3$  (b)  $NH_3$ (c)  $P_4O_6$  (d)  $P_4O_{10}$  (1991)

#### 7.8 Phosphorus Halides

24. Identify the incorrect statement related to PCI from

the following :

- (a)  $PCl_5$  molecule is non-reactive.
- (b) Three equatorial P Cl bonds make an angle of 120° with each other.
- (c) Two axial P Cl bonds make an angle of  $180^{\circ}$  with each other.
- (d) Axial P Cl bonds are longer than equatorial P – Cl bonds. (NEET 2019)

- **25.**  $PCl_3$  reacts with water to form
  - (a) PH<sub>3</sub>
    (b) H<sub>3</sub>PO<sub>3</sub>, HCl
    (c) POCl<sub>3</sub>
    (d) H<sub>3</sub>PO<sub>4</sub>
    (1991)

# 7.9 Oxoacids of Phosphorus

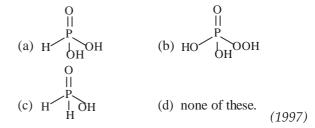
- **26.** Which of the following oxoacids of phosphorus has strongest reducing property?
  - (a)  $H_4P_2O_7$  (b)  $H_3PO_3$
  - (c)  $H_3PO_2$  (d)  $H_3PO_4$

(Odisha NEET 2019)

- 27. Which is the correct statement for the given acids?
  - (a) Phosphinic acid is a monoprotic acid while phosphonic acid is a diprotic acid.
  - (b) Phosphinic acid is a diprotic acid while phosphonic acid is a monoprotic acid.
  - (c) Both are diprotic acids.
  - (d) Both are triprotic acids. (NEET-I 2016)
- **28.** Strong reducing behaviour of  $H_3PO_2$  is due to
  - (a) high electron gain enthalpy of phosphorus
  - (b) high oxidation state of phosphorus
  - (c) presence of two —OH groups and one P—H bond
  - (d) presence of one —OH group and two P—H bonds. (2015)
- **29.** Which of the following statements is not valid for oxoacids of phosphorus?
  - (a) Orthophosphoric acid is used in the manufacture of triple superphosphate.
  - (b) Hypophosphorous acid is a diprotic acid.
  - (c) All oxoacids contain tetrahedral four coordinated phosphorus.
  - (d) All oxoacids contain atleast one P\_O unit and one P-OH group. (2012)
- **30.** Oxidation states of P in H<sub>4</sub>P<sub>2</sub>O<sub>5</sub>, H<sub>4</sub>P<sub>2</sub>O<sub>6</sub>, H<sub>4</sub>P<sub>2</sub>O<sub>7</sub> are respectively

(a) 
$$+3, +5, +4$$
 (b)  $+5, +3, +4$   
(c)  $+5, +4, +3$  (d)  $+3, +4, +5$  (2010)

- **31.** How many bridging oxygen atoms are present in  $P_4O_{10}$ ?
  - (a) 6 (b) 4 (c) 2 (d) 5 (*Mains* 2010)
- 32. The structural formula of hypophosphorous acidis



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33.	H <sub>3</sub> PO <sub>2</sub> is the molecular phosphorus. Its name an (a) phosphorous acid a (b) hypophosphorous a (c) hypophosphorous a (d) hypophosphoric acid	nd basicity respectiv nd two cid and two cid and one	
34.	Which one of the follow laboratory for fast dryin (a) Phosphorus pentoxida (c) Anhydrous calcium (d) Na <sub>3</sub> PO <sub>4</sub>	ng of neutral gases? e (b) Active charcoa	
35.	P <sub>2</sub> O <sub>5</sub> is heated with wat (a) hypophosphorous a (b) phosphorous acid (d) orthophosphoric acid	cid (c) hypophosphor	ic acid (1991)
36.	Basicity of orthophosph (a) 2 (c) 4	noric acid is (b) 3 (d) 5	(1991)
37.	<ul><li>When orthophosphoric product formed is</li><li>(a) PH<sub>3</sub></li><li>(c) H<sub>3</sub>PO<sub>3</sub></li></ul>	acid is heated to 60 (b) P <sub>2</sub> O <sub>5</sub> (d) HPO <sub>3</sub>	0°C, the (1989)
7.	0 Group 16 Elemer	nts	
38.	Which is the correct the $(E = O, S, Se, Te and Pc)$ (a) $H_2Se < H_2Te < H_2Pc$ (b) $H_2S < H_2O < H_2Se$ (c) $H_2O < H_2S < H_2Se$ (d) $H_2Po < H_2Te < H_2Se$	o) ? o < H <sub>2</sub> O < H <sub>2</sub> S < H <sub>2</sub> Te <h<sub>2Po &lt; H<sub>2</sub>Te <h<sub>2Po</h<sub></h<sub>	
39.	Acidity of diprotic acid	s in aqueous solutio	ons

- 39. Acidity of diprotic acids in aqueous solutions increases in the order
  (a) H<sub>2</sub>S < H<sub>2</sub>Se < H<sub>2</sub>Te (b) H<sub>2</sub>Se < H<sub>2</sub>S < H<sub>2</sub>Te
  - (a)  $\Pi_2 S < \Pi_2 S e < \Pi_2 Ie(0) \Pi_2 S e < \Pi_2 S < \Pi_2 Ie(0) \Pi_2 S e < \Pi_2 S e < \Pi_2 Ie(0) II S e < I$
  - (c)  $H_2Te < H_2S < H_2Se$  (d)  $H_2Se < H_2Te < H_2Se$

(2014)

40. Which of the following bonds has the highest energy?
(a) S–S
(b) O–O
(c) Se–Se
(d) Te–Te
(1996)

# 7.11 Dioxygen

- **41.** Which of the following does not give oxygen on heating?
  - (a)  $K_2Cr_2O_7$  (b)  $(NH_4)_2Cr_2O_7$
  - (c)  $KClO_3$  (d)  $Zn(ClO_3)_2$

(NEET 2013)

- **42.** Which would quickly absorb oxygen?
  - (a) Alkaline solution of pyrogallol
  - (b) Conc. H<sub>2</sub>SO<sub>4</sub>
  - (c) Lime water
  - (d) Alkaline solution of CuSO<sub>4</sub> (1991)

- **43.** Oxygen will directly react with each of the following elements except
  - (a) P (b) Cl (c) Na (d) S (1989)
- **44.** It is possible to obtain oxygen from air by fractional distillation because
  - (a) oxygen is in a different group of the periodic table from nitrogen
  - (b) oxygen is more reactive than nitrogen
  - (c) oxygen has higher b.pt. than nitrogen
  - (d) oxygen has a lower density than nitrogen. (1989)

# 7.12 Simple Oxides

**45.** Match the following :

		0		
Oxid	le		Nat	ure
(A) CO			(i) Basic	2
(B) BaO			(ii) Neu	tral
$(C) Al_2O_2$	3		(iii) Acid	lic
$(D) Cl_2O_2$	7		(iv) Amp	ohoteric
Which of	the fol	lowing	is correct	option?
(A)	<b>(B)</b>	(C)	(D)	
(a) (i)	(ii)	(iii)	(iv)	
(b) (ii)	(i)	(iv)	(iii)	
(c) (iii)	(iv)	(i)	(ii)	
(d) (iv)	(iii)	(ii)	(i)	(NEET 2020)

# 7.13 Ozone

- **46.** The angular shape of ozone molecule  $(O_3)$  consists of
  - (a)  $1\sigma$  and  $1\pi$  bond (b)  $2\sigma$  and  $1\pi$  bond
  - (c)  $1\sigma$  and  $2\pi$  bonds (d)  $2\sigma$  and  $2\pi$  bonds.(2008)
- 47. The gases respectively absorbed by alkaline pyrogallol and oil of cinnamon are
  (a) O<sub>3</sub>, CH<sub>4</sub>
  (b) O<sub>2</sub>, O<sub>3</sub>
  - (c)  $SO_2$ ,  $CH_4$  (d)  $N_2O$ ,  $O_3$  (1989)

# 7.15 Sulphur Dioxide

- **48.** Nitrogen dioxide and sulphur dioxide have some properties in common. Which property is shown by one of these compounds, but not by the other?
  - (a) Is soluble in water.
  - (b) Is used as a food preservative.
  - (c) Forms 'acid-rain'.
  - (d) Is a reducing agent. (2015, Cancelled)
- **49.** Sulphur trioxide can be obtained by which of the following reaction?
  - (a)  $CaSO_4 + C \xrightarrow{\Delta}$

(b) 
$$\operatorname{Fe}_2(\operatorname{SO}_4)_3 \xrightarrow{\Delta}$$

(c) 
$$S + H_2SO_4 \xrightarrow{\Delta} \Delta$$

(d)  $H_2SO_4 + PCl_5$  —

(2012)

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7.16 Oxoacids of Sulphur 50. Which of the following oxoacid of sulphur has -O - O - linkage(a)  $H_2SO_3$ , sulphurous acid (b)  $H_2SO_4$ , sulphuric acid (c)  $H_2S_2O_8$ , peroxodisulphuric acid (d)  $H_2S_2O_7$ , pyrosulphuric acid (NEET 2020) 51. Identify the correct formula of oleum from the following : (a)  $H_2S_2O_7$ (b)  $H_2SO_3$ (c)  $H_2SO_4$ (d)  $H_2S_2O_8$ (OdishaNEET2019) 52. In which pair of ions both the species contain S - Sbond? (a) S O  $^{2-}$ , S O $^{2-}$ (b) S  $O^{2-}$ , S  $O^{2-}$ (d)  $S_{2}^{2}O_{7}^{\frac{7}{2}}, S_{2}^{2}O_{3}^{\frac{8}{2}}$ (c)  $S^4O^{2-}, S^2O^{3-}$ 6 (NEET 2017) 53. Oleum is

(a) castor oil
(b) oil of vitriol
(c) fuming H<sub>2</sub>SO<sub>4</sub>
(d) none of these. (1991)

# 7.17 Sulphuric Acid

**54.** Match List I (substances) with List II (processes) employed in the manufacture of the substances and select the correct option.

server and control option		
List I	List II	
(Substances)	(Proces	sses)
(A) Sulphuric acid	(i) Haber's	process
(B) Steel	(ii) Bessem	er's
	process	
(C) Sodium	(iii)Lebland	e process
hydroxide		-
(D) Ammonia	(iv) Contact	t process
(a) A - (i), B - (iv), C -	(ii), D - (iii)	
(b) A - (i), B - (ii), C - (	(iii), D -(iv)	
(c) A - (iv), B - (iii), C	- (ii), D -(i)	
(d) A - (iv), B - (ii), C -	· (iii), D- (i)	(Mains 2010)

# 7.18 Group 17 Elements

- 55. Which of the following statements is not true for halogens?
  - (a) All form monobasic oxyacids.
  - (b) All are oxidizing agents.
  - (c) All but fluorine show positive oxidation states.
  - (d) Chlorine has the highest electron-gain (*NEET 2018*)
- **56.** Which one of the following orders is correct for the bond dissociation enthalpy of halogen molecules?

(a)  $Br_2 > I_2 > F_2 > Cl_2$  (b)  $F_2 > Cl_2 > Br_2 > I_2$ (c)  $I_2 > Br_2 > Cl_2 > F_2$  (d)  $Cl_2 > Br_2 > F_2 > I_2$ (*NEET-I* 2016)

- 57. The variation of the boiling points of the hydrogen halides is in the order HF > HI > HBr > HCl.What explains the higher boiling point of hydrogen fluoride?
  - (a) There is strong hydrogen bonding between HF molecules.
  - (b) The bond energy of HF molecules is greater than in other hydrogen halides.
  - (c) The effect of nuclear shielding is much reduced in fluorine which polarises the HF molecule.
  - (*d*) The electronegativity of fluorine is much higher than for other elements in the group. (2015)
- 58. Among the following which is the strongest oxidising agent? (b) I

(c) 
$$Cl^{2}_{2}$$
 (d)  $\vec{F}_{2}$  (2009)

- **59.** Which one of the following arrangements does not give the correct picture of the trends indicated against it?
  - (a)  $F_2 > Cl_2 > Br_2 > I_2$ : Bond dissociation energy
  - (b)  $F_2 > Cl_2 > Br_2 > I_2$ : Electronegativity
  - (c)  $F_2 > Cl_2 > Br_2 > I_2$ : Oxidizing power
  - (d)  $F_2>Cl_2>Br_2>I_2$  : Electron gain enthalpy  $(2008) \label{eq:result}$
- **60.** Which one of the following orders is not in accordance with the property stated against it?
  - (a)  $F_2 > Cl_2 > Br_2 > I_2$ : Bond dissociation energy (b)  $F_2 > Cl_2 > Br_2 > I_2$ : Oxidising power
  - (c) HI > HBr > HCl > HF: Acidic property in water
  - (d)  $F_2 > Cl_2 > Br_2 > I_2$ : Electronegativity (2006)
- **61.** Which statement is wrong?
  - (a) Bond energy of  $F_2 > Cl_2$
  - (b) Electronegativity of F > Cl
  - (c) F is more oxidising than Cl
  - (d) Electron affinity of Cl > F (2000)
- **62.** Which of the following has the greatest electron affinity?
  - (a) I (b) Br (c) F (d) Cl (1996)
- **63.** Which of the following displaces Br<sub>2</sub> from an aqueous solution containing bromide ions?
  - (a)  $I_2$  (b)  $I_3^-$ (c)  $Cl_2$  (d) Cl (1994)

(1993)

**64.** Which of the following species has four lone pairs of electrons?

(a) I	(b) O
(c) Cl <sup>-</sup>	(d) He

# 7.19 Chlorine

#### 65. Match the following: (A) Pure nitrogen (i) Chlorine (B) Haber process (ii) Sulphuricacid (C) Contact process (iii) Ammonia (D) Deacon's process (iv) Sodium azide or Barium azide Which of the following is the correct option? (A) **(B)** (C) (D) (a) (iv) (iii) (ii) (i) (b) (i) (ii) (iii) (iv) (c) (ii) (iv) (i) (iii) (iv) (ii) (NEET 2019) (d) (iii) (i) 66. When Cl<sub>2</sub> gas reacts with hot and concentrated sodium hydroxide solution, the oxidation number of chloring changes from -5 (b) zero to -1 and zero to +5(c) zero to -1 and zero to +3(d) zero to +1 and zero to -367. Which of the following is used in the preparation of chlorine? (a) Both $MnO_2$ and $KMnO_4$

(b) Only KMnO<sub>4</sub>

(d) Either  $MnO_2$  or  $KMnO_4$ 

(c) Only MnO<sub>2</sub>

(1999)

(2012)

- 68. Which of the following elements is extracted commercially by the electrolysis of an aqueous solution of its compound?
  - (a) Cl (b) Br (c) Al (d) Na (1993)
- 69. When chlorine is passed over dry slaked lime at room temperature, the main reaction product is
  - (a)  $Ca(ClO_2)_2$ (b) CaCl<sub>2</sub>
  - (c) CaOCl<sub>2</sub> (d) Ca(OCl)<sub>2</sub> (1992)
- 70. In the manufacture of bromine from sea water, the mother liquor containing bromides is treated with (a) carbon dioxide (b) chlorine
  - (c) iodine (d) sulphur dioxide.

(1992)

- **71.** The bleaching action of chlorine is due to (a) reduction (b) hydrogenation
  - (c) chlorination (d) oxidation. (1991)
- 7.20 Hydrogen Chloride
- 72. Bleaching powder reacts with a few drops of conc. HCl to give
  - (a) chlorine (b) hypochlorous acid
  - (c) calcium oxide (d) oxygen. (1989)

# 7.21 Oxoacids of Halogens

- 73. Among the following, the correct order of acidity is
  - (a)  $HClO_2 < HClO < HClO_3 < HClO_4$
  - (b)  $HClO_4 < HClO_2 < HClO < HClO_3$
  - (c)  $HClO_3 < HClO_4 < HClO_2 < HClO$
  - (d)  $HClO < HClO_2 < HClO_3 < HClO_4$

(NEET-I 2016, 2007, 2005)

- 74. Which of the statements given below is incorrect? (a)  $O_3$  molecule is bent.
  - (b) ONF is isoelectronic with  $O_2N$ .
  - (c)  $OF_2$  is an oxide of fluorine.
  - (d)  $Cl_2O_7$  is an anhydride of perchloric acid.

(2015)

75. The correct order of increasing bond angles in the following species is

(a) 
$$Cl_{2}O < ClO_{2} < ClO_{2}$$
  
(b)  $ClO_{2} < Cl_{2}O < ClO_{2}^{-}$   
(c)  $Cl_{2}O < ClO_{2}^{-} < ClO_{2}^{-}$   
(d)  $ClO_{2}^{-} < ClO_{2}^{-} < ClO_{2}^{-}$   
(2010)

76. Which one of the following oxides is expected to exhibit paramagnetic behaviour?

(a) CO<sub>2</sub> (b)  $SiO_2$ 

(c)  $SO_2$ (d)  $ClO_2$ (2005)

# 7.22 Interhalogen Compounds

77. Match the interhalogen compounds of column-I with the geometry in column-II and assign the correct code.

(	Colı	ımn I		Column II	
(A) 2	XX'		(i) '	T-shape	
(B) 2	$XX'_3$		(ii) Pentagonal bipyramidal		
(C) 2	$XX'_5$		(iii)	Linear	
(D) 2	$XX'_7$		(iv)	Square pyramic	lal
			(v) '	Tetrahedral	
Cod	le:				
	Α	B	С	D	
(a) (	(iii)	(i)	(iv)	(ii)	
(b) (	(v)	(iv)	(iii)	(ii)	
(c) (	(iv)	(iii)	(ii)	(i)	
(d) (	(iii)	(iv)	(i)	(ii)	(NEET 2017)

# 7.23 Group-18 elements

78. Match the Xenon compounds in Column-I with its structure in Column-II and assign the correct code.

Co	Column-I		Column-II
(A)	XeF <sub>4</sub>	(i)	pyramidal
(B)	XeF <sub>6</sub>	(ii)	square planar
(C)	XeOF <sub>4</sub>	(iii)	distorted octahedral
(D)	XeO <sub>3</sub>	(iv)	square pyramidal

#### The p-Block Elements

	(A)	<b>(B)</b>	(C)	(D)	(a) XeO <sub>4</sub>
(	(a) (iii)	(iv)	(i)	(ii)	(b) There
(	(b) (i)	(ii)	(iii)	(iv)	(c) There
(	(c) (ii)	(iii)	(iv)	(i)	(d) XeO <sub>4</sub>
(	(d) (ii)	(iii)	(i)	(iv)	
				(NEET 2019, NEET-I 2016)	80 Which cor

**79.** Identify the incorrect statement, regarding the molecule XeO<sub>4</sub>.

(b) There are four <i>p</i>	$p\pi - d\pi$ bonds.			
(c) There are four $sp^3 - p$ , $\sigma$ bonds.				
(d) XeO <sub>4</sub> molecule is tetrahedral.				
	(Karnataka	NEET 2013)		
80. Which compound ha	as planar structure?			
(a) $XeF_4$	(b) XeOF <sub>2</sub>			
(c) $XeO_2F_2$	(d) XeO <sub>4</sub>	(2000)		
	<ul> <li>(b) There are four p</li> <li>(c) There are four s</li> <li>(d) XeO<sub>4</sub> molecule</li> <li>80. Which compound has</li> <li>(a) XeF<sub>4</sub></li> </ul>	( <i>Karnataka</i> 80. Which compound has planar structure? (a) XeF <sub>4</sub> (b) XeOF <sub>2</sub>		

	ANSWER KEY																		
1.	(c)	2.	(b)	3.	(c)	4.	(c)	5.	(a)	6.	(a)	7.	(a)	8.	(b)	9.	(a)	10.	(d)
11.	(d)	12.	(b)	13.	(b)	14.	(d)	15.	(a)	16.	(c)	17.	(c)	18.	(d)	19.	(b)	20.	(b)
21.	(a)	22.	(a)	23.	(a)	24.	(a)	25.	(b)	26.	(c)	27.	(a)	28.	(d)	29.	(b)	30.	(d)
31.	(a)	32.	(c)	33.	(c)	34.	(a)	35.	(d)	36.	(b)	37.	(d)	38.	(d)	39.	(a)	40.	(a)
41.	(b)	42.	(a)	43.	(b)	44.	(c)	45.	(b)	46.	(b)	47.	(b)	48.	(b)	49.	(b)	50.	(c)
51.	(a)	52.	(a)	53.	(c)	54.	(d)	55.	(c)	56.	(d)	57.	(a)	58.	(d)	59.	(a,d)	60.	(a)
61.	(a)	62.	(d)	63.	(c)	64.	(c)	65.	(a)	66.	(b)	67.	(a)	68.	(a)	69.	(c)	70.	(b)
71.	(d)	72.	(a)	73.	(d)	74.	(c)	75.	(d)	76.	(d)	77.	(a)	78.	(c)	79.	(a)	80.	(a)

# **Hints & Explanations**

1. (c) :  $N_2H_4 \Rightarrow 2x + 4(+1) = 0$   $\Rightarrow 2x + 4 = 0$   $\Rightarrow x = -2$   $NH_3 \Rightarrow x + 3(+1) = 0 \Rightarrow x = -3$   $N_3H \Rightarrow 3x + 1(+1) = 0$   $\Rightarrow 3x + 1 = 0 \Rightarrow x = -1/3$   $NH_2OH \Rightarrow x + 2 + 1(-2) + 1 = 0$   $\Rightarrow x + 1 = 0 \Rightarrow x = -1$ Thus, highest oxidation state is -1/3.

2. (b) : For strong  $\pi$ -bonding,  $p\pi - p\pi$  bonding should be strong. In case of P, due to larger size as compared to N-atom,  $p\pi - p\pi$  bonding is not sostrong.

**3.** (c) : Among N, P, As and Sb, the former has highest electronegativity (*EN*) so its oxide is most acidic.

As the electronegativity value of element increases, the acidic character of the oxide also increases.

4. (c) : Due to greater electronegativity of nitrogen, dipole moment for  $NH_3$  is greater.

5. (a) : All the hydrides of group V elements have one lone pair of electrons on their central atom. Therefore, they can act as Lewis bases. The basic character of these hydrides decreases down the group.

6. (a) : The acidic character of the oxides decreases with the decrease in the oxidation state and also decreases

down the group.

7. (a) : Nitrogen cannot form pentahalides because it cannot expand its octet due to non-availability of d-orbitals.

8. (b) : Boiling point of hydrides increases with increase in atomic number but ammonia has exceptionally high boiling point due to hydrogen bonding. Thus, the correct order of boiling point is,

 $BiH_3 > SbH_3 > NH_3 > AsH_3 > PH_3$ 

9. (a) : Nitrogen molecule is diatomic containing a triple bond between two N atoms,  $N \equiv N$  therefore, nitrogen molecule is formed by sharing six electrons.

10. (d) :  $N_2$  molecule contains triple bond between N atoms having very high dissociation energy (946 kJ mol<sup>-1</sup>) due to which it is relatively inactive.

11. (d) : NH<sub>4</sub>Cl + NaNO<sub>2</sub><sup>Heat</sup> NH<sub>4</sub>NO<sub>2</sub> + NaCl  

$$\downarrow^{\Lambda}$$
  
N<sub>2</sub> + 2H<sub>2</sub>O

**12.** (b) : In case of nitrogen, *d*-orbitals are not available for bonding. N :  $1s^2 2s^2 2p^3$ 

13. (b): 
$$NH_2 - \overset{O}{\overset{\parallel}{C}} - NH_2 + H_2O \longrightarrow NH_2COONH_4$$
  
Urea (A)  
 $\longrightarrow 2NH_3 + CO_2$   
(B)  
 $Cu^{2+}_{(aq)} + 4NH_3 \longrightarrow [Cu(NH_3)_4]^{2+}_{(dep blue)}$   
(C)

14. (d) : Aqueous solution of ammonia contains NH<sup>+</sup> and OH<sup>-</sup> ions.

$$\frac{\mathrm{NH} + \mathrm{H} \mathrm{O} \mathrm{NH}^{+} + \mathrm{OH}^{-}}{3 2} 4$$

**15.** (a) :  $NO_2$  is paramagnetic due to the presence of one unpaired electron.

**16.** (c) : When two molecules of nitric acid undergoes heating, loose a water molecule to form ananhydride.

 $HONO_2 \xrightarrow{\Lambda} N_2O_5 + H_2O$ 

Thus, N<sub>2</sub>O<sub>5</sub> is nitric acid anhydride. 17. (c) :  $Cu + 4HNO_3(conc.) \longrightarrow Cu(NO_3)_2 + 2NO_2$ + 2HO

18. (d) : Zinc is on the top position of hydrogen in electrochemical series. So, Zn displaces H<sub>2</sub> from dilute H<sub>2</sub>SO<sub>4</sub> and HCl with liberation of H<sub>2</sub>.

$$Zn + H_2SO_4 = 2nSO_4 + H_2$$

On the other hand, HNO<sub>3</sub> is one oxidising agent. Hydrogen obtained in the reaction is converted into  $H_2O$ .

$$Zn + 2HNO_3 Zn(NO_3)_2 + 2H$$
$$2HNO_3 \longrightarrow H_2O + 2NO_2 + O$$
$$2H + O \longrightarrow H_2O$$

**19. (b)** :  $C_{12}H_{22}O_{11} + 18[O] \longrightarrow 6(COOH)_2 + 5H_2O$ Cane sugar From HNO<sub>3</sub> Oxalic acid

20. (b): White phosphorus has low ignition temperature so it is most reactive among all the allotropes.

**21.** (a) : Red phosphorus is insoluble in  $CS_2$  and only white P is soluble in CS<sub>2</sub>.

22. (a) : 
$$Ca_3P_2 + 6H_2O \rightarrow 3Ca(OH)_2 + 2PH_{3(g)}$$
  
(X) (Y)  
(Rotten  
fish smell)

 $3CuSO_4 + 2PH_3 \longrightarrow Cu_3P_2 + 3H_2SO_4$ (Copper (Y)phosphide)

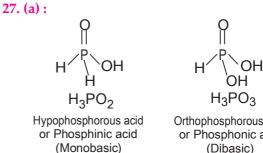
23. (a) : 
$$PH_4I + NaOH \rightarrow NaI + PH_3 + H_2O$$

**24.** (a) : It is a reactive gas as it easily provides  $Cl_2$  gas.

240 pm 
$$\downarrow$$
 P  $Cl$  202 pm  $Cl$ 

**26.** (c) : Acids which contain P - H bonds have strong reducing properties. Among the given compounds,  $H_3PO_2$  is the strongest reducing agent as it contains two P – H bonds. 0





Orthophosphorous acid or Phosphonic acid (Dibasic)

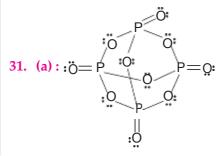
28. (d) : All oxyacids of phosphorus which have P—H bonds act as strong reducing agents. H<sub>3</sub>PO<sub>2</sub> has two P-H bonds hence, it acts as a strong reducing agent.



29. (b) : Hypophosphorous acid is a monoprotic acid.

**30.** (d) : The oxidation state can be calculated as :

 $H_4P_2O_5: +4 + 2x + 5(-2) = 0 \Longrightarrow 2x - 6 = 0 \Longrightarrow x = +3$  $H_4P_2O_6$ : +4 + 2x + 6(-2) = 0  $\Rightarrow$  2x - 8 = 0  $\Rightarrow$  x = +4  $H_4P_2O_7: +4 + 2x + 7(-2) = 0 \implies 2x - 10 = 0 \implies x = +5$ 



32. (c) : The formula of hypophosphorous acid is  $H_3PO_2$  as shown in (c). It is a monobasic acid.

**33.** (c) :  $H_3PO_2$  is named as hypophosphorous acid. As it contains only one P-OH group, its basicity is one.

34. (a) :  $P_2O_5$  absorbs moisture much readily than anhydrous CaCl<sub>2</sub>.

**35. (d)** :  $P_2O_5 + 3H_2O \xrightarrow{\Delta} 2H_3PO_4$ 

**36.** (b) : Orthophosphoric acid, H<sub>3</sub>PO<sub>4</sub> contains three P-OH groups and is therefore, tribasic.

**37.** (d) : On heating, it gives pyrophosphoric acid at 525 K and metaphosphoric acid at 875 K.

$$\begin{array}{cccc} 2H & PO \\ & & 525 \text{ K} \\ & & -H_2O \end{array} \rightarrow H \begin{array}{c} P & O \\ & & +I_2O \end{array} \xrightarrow{875 \text{ K}} -H_2O \end{array} \xrightarrow{875 \text{ K}} 2HPO_3 \\ \hline \\ Orthophosphoric \\ acid \end{array} \xrightarrow{\text{Netaphosphoric}} \begin{array}{c} \text{Metaphosphoric} \\ acid \end{array} \xrightarrow{\text{Retarms}} acid \end{array}$$

**38.** (d): The thermal stability of hydrides decreases from H<sub>2</sub>O to H<sub>2</sub>Po. This is because as the size of atom

*E* in  $H_2E$  increases, the bond  $H_-E$  becomes weaker and thus, breaks on heating. Therefore, the correct order of thermal stability is  $H_2Po < H_2Te < H_2Se < H_2S < H_2O$ .

**39.** (a) : As the atomic size increases down the group, the bond length increases and the bond strength decreases and the cleavage of E-H bond becomes easier thus, more will be the acidity. Thus, the correct order is :  $H_2S < H_2Se < H_2Te$ .

40. (a) : Bond energy of S - S is exceptionally high due to its catenation tendency.

B.E. O—O S—S Se— Se Te— Te  
(kJ mol<sup>-1</sup>): 142 226 172 126  
**41.** (b): (NH<sub>4</sub>)<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>
$$\xrightarrow{\Delta}$$
 N<sub>2</sub>+Cr<sub>2</sub>O<sub>3</sub>+4H<sub>2</sub>O

$$Zn(ClO_3)_2 \xrightarrow{\Lambda} ZnCl_2 + 3O_2$$

 $\text{KClO}_3 \xrightarrow{\Delta} \text{KCl} + 3/2\text{O}_2$ 

 $2K_2Cr_2O_7 \xrightarrow{\Delta} 2K_2CrO_4 + Cr_2O_3 + 3/2O_2$ 

**42.** (a) : Alkaline solution of pyrogallol absorbs oxygen quickly.

**43.** (b) : Chlorine does not react directly with oxygen.

44. (c) : Air is liquefied by making use of the Joule

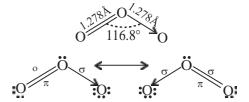
- Thomson effect (cooling by expansion of the gas).

Water vapour and  $CO_2$  are removed by solidification. The remaining constituents of liquid air *i.e.*, liquid oxygen and liquid nitrogen are separated by means of fractional distillation as fractional distillation is a process of separation of mixture based on the difference in their boiling points. (b.pt. of  $O_2 = -183^{\circ}C$  : b.pt. of  $N_2 = -195.8^{\circ}C$ ).

**45.** (b) : CO — neutral, BaO — basic,

 $Al_2O_3$  — amphoteric and  $Cl_2O_7$  — acidic.

**46.** (b): The angular shape of ozone molecule  $(O_3)$ :



**47.** (b) : Alkaline pyrogallol absorbs O<sub>2</sub> and oil of cinnamon absorbs O<sub>3</sub>.

48. (b) : NO<sub>2</sub> is not used as a foodpreservative. 49. (b) : Fe (SO)  $\xrightarrow{\Delta}$  Fe O + 3SO  $\xrightarrow{2}$   $\xrightarrow{43}$   $\xrightarrow{3}$   $\xrightarrow{3}$ 

50. (c) : Peroxodisulphuric acid,  $H_2S_2O_8$  has -O - O - linkage.

$$O_{OH} = O_{OH} = O$$

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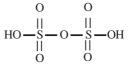
 $(S_{2}Q^{2})$ 

51. (a)

 $(S_2Q^{2-})$ 

53. (c) : Pyrosulphuric acid or oleum (+6) is H S O

which is obtained by dissolving SO<sub>3</sub> and is called fuming sulphuric acid.



#### 54. (d)

55. (c) : All halogens show both positive and negative oxidation states while fluorine shows only negative oxidation state except +1 in HOF.

56. (d): The order of bond dissociation enthalpy is :

B.E. (in kJ mol<sup>-1</sup>) 242.6 192.8 158.8 151.1 A reason for this anomaly is the relatively large electronelectron repulsion among the lone pairs in  $F_2$  molecule where they are much closer to each other than in case of Cl<sub>2</sub>.

57. (a) : HF forms strong intermolecular H-bonding

due to high electronegativity of F. Hence, the boiling point of HF is abnormally high. Boiling points of other hydrogen halides gradually increase from HCl to HI due to increase in size of halogen atoms from Cl to I which further increase the magnitude of van der Waals' forces.

**58.** (d) : Standard reduction potentials of halogens are positive and decrease from fluorine to iodine. So,  $F_2$  is the strongest oxidising agent.

**59.** (a, d): In case of diatomic molecules  $(X_2)$  of halogens the bond dissociation energy decreases in the order :  $Cl_2 > Br_2 > F_2 > I_2$ . This is due to relatively large electron-electron repulsion among the lone pairs is  $F_2$  than in case of  $Cl_2$ .

The oxidising power, electronegativity and reactivity decrease in the order :  $F_2 > Cl_2 > Br_2 > I_2$ 

Electron gain enthalpy of halogens follows the given order :

 $Cl_2 > F_2 > Br_2 > I_2$ 

The low value of electron gain enthalpy of fluorine is probably due to small size of fluorine atom.

**60.** (a) : X - X bond F - F Cl - Cl Br - Br I - I Bond dissociation 38 57 45.5 35.6 energy (kcal/mol)

The lower value of bond dissociation energy of fluorine is due to the high inter-electronic repulsions between non-bonding electrons in the 2p-orbitals of fluorine. As a result F – F bond is weaker in comparison to Cl – Cl and Br – Br bonds.

61. (a): Due to more repulsion in between non-

bonding electron pairs (2p) of two fluorines (due to small size of F-atom) in comparison to non-bonding electron pairs (3p) in chlorine, the bond energy of F<sub>2</sub> is less than Cl<sub>2</sub>.

*B.E.* (F<sub>2</sub>) = 158.8 kJ/mole and *B.E.* (Cl<sub>2</sub>) = 242.6 kJ/mole

**62.** (d) : In general, the electron affinity decreases from top to bottom in a group. But in group 17, fluorine has lower electron affinity as compared to chlorine due to very small size of fluorine atom. As a result, there are strong interelectronic repulsions in the relatively small 2*s* orbitals of fluorine and thus, the incoming electron does not experience much attraction.

**63.** (c) : Since chlorine is stronger oxidising agent than bromine, therefore it will displace bromine from an aqueous solution containing bromide ions.

$$Cl_2 + 2Br^- \rightarrow 2Cl^- + Br_2$$

64. (c) : Outer electronic configuration of Cl  
= 
$$3s^23p^23p^23p^23p^1$$
  
Outer electronic configuration of Cl<sup>-</sup>  
=  $3s^23p^23p^23p^2$ , *i.e.*, 4 lone pair of electrons  
65. (a)  
66. (b):  $3Cl_2 + 6NaOH \rightarrow 5NaCl + NaClO_3 + 3H_2O$ 

This is an example of disproportionation reaction and oxidation state of chlorine changes from 0 to -1 and +5.

67. (a) : 
$$MnO_2 + 4HCl \rightarrow MnCl_2 + 2H_2O + Cl_2\uparrow$$
  
2KMnO<sub>4</sub> + 16HCl  $\rightarrow$  2KCl + 2MnCl<sub>2</sub> + 8H<sub>2</sub>O + 5Cl<sub>2</sub>↑

**68.** (a) : Chlorine is obtained by the electrolysis of brine (concentrated NaCl solution). Chlorine is liberated at anode.

69. (c) : Ca(OH)<sub>2</sub> + Cl<sub>2</sub>  $\rightarrow$  CaOCl<sub>2</sub> + H<sub>2</sub>O

**70.** (b) : Bromide in the mother liquor (containing  $MgBr_2$ ) is oxidised to  $Br_2$  by passing  $Cl_2$  which is a stronger oxidising agent.

 $2Br^-\!+Cl_2 \to Br_2+2Cl^-$ 

71. (d) : Bleaching action of chlorine is due to oxidation in presence of moisture. Bleaching effect is permanent.

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

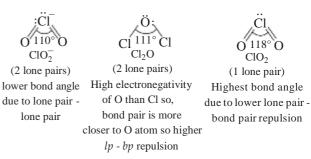
Colouring matter +  $[O] \rightarrow$  Colourless matter

72. (a) :  $CaOCl_2 + 2HCl \rightarrow CaCl_2 + H_2O + Cl_2$ 

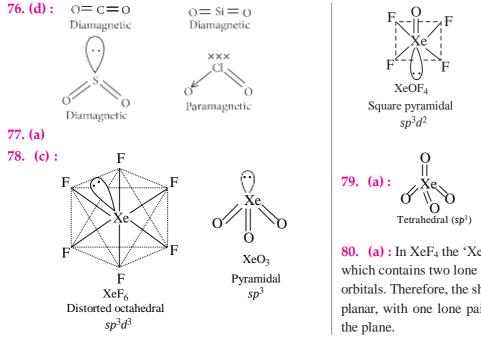
73. (d) : The acidic character of the oxoacids increases with increase in oxidation number of the halogen atom *i.e.*,  $HCIO < HCIO_2 < HCIO_3 < HCIO_4$ .

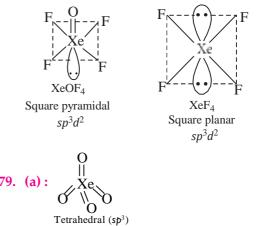
This can be explained on the basis of relative stability of the anions left after removal of a proton. Since the stability of the anion decreases in the order :  $\text{ClO}_4^- > \text{ClO}_3^- > \text{ClO}_2^- > \text{ClO}^-$ , acid strength also decreases in the same order.

74. (c) :  $OF_2$  (oxygen difluoride) is a fluoride of oxygen because fluorine is more electronegative than oxygen. 75. (d) :



#### The p-Block Elements





80. (a) : In XeF<sub>4</sub> the 'Xe' atom is  $sp^3d^2$  hybridised, which contains two lone pair orbitals and four bond pair orbitals. Therefore, the shape of XeF<sub>4</sub> molecule is square planar, with one lone pair orbital over and other below