

Objective Questions

#### Alkali metals

- 1. As compared to potassium, sodium has[MP PMT 1985]
  - (a) Lower electronegativity
  - (b) Higher ionization potential
  - (c) Greater atomic radius
  - (d) Lower melting point
- Potassium is kept in 2.

[CPMT 1976]

- (a) Alcohol
- (b) Water
- (c) Kerosene
- (d) Liquid ammonia
- The product obtained on fusion of  $BaSO_4$  and 3.  $Na_2CO_3$  is

[AFMC 2005]

- (a)  $BaCO_3$
- (b) BaO
- (c)  $Ba(OH)_2$
- (d) BaHSO A
- Which of the following statement is correct 4. regarding alkali metals [NCERT 1981]
  - (a) Cation is less stable than the atom
  - (b) Cation is smaller than the atom
  - (c) Size of cation and atom is the same
  - (d) Cation is greater in size than the atom
- Valency electrons in alkali metals are [CPMT 1972] 5.
  - (a) 1

(b) 7

(c) 4

- (d) 2
- Magnitude of which of the following property of 6. increases with the increase of alkali metals atomic number

[MP PMT 1987]

- (a) Electronegativity
- (b) Ionic radius
- (c) First ionization energy (d)
- Melting point
- 7. As compared to lithium, sodium reacts quickly with water because [NCERT 1978, 80]
  - (a) Its molecular weight is less
  - (b) It is stronger electronegative
  - (c) It is stronger electropositive
  - (d) It is a metal
- 8. Which is an ore of potassium

[DPMT 1984; CPMT 1986; Kurukshetra CEE 1998]

- (a) Carnellite
- (b) Cryolite
- (c) Bauxite
- (d) Dolomite
- $Na_2CO_3$  can be manufactured by Solvey's process 9. 18. but  $K_2CO_3$  cannot be prepared because
  - (a)  $K_2CO_3$  is more soluble

- (b)  $K_2CO_3$  is less soluble
- (c)  $KHCO_3$  is more soluble than  $NaHCO_3$
- (d) KHCO<sub>2</sub> is less soluble than NaHCO<sub>2</sub>
- Which of the following alkali metals is smallest in

[CPMT 1990]

- (a) *Rb*
- (b) K
- (c) Na
- (d) Li
- 11. When potassium dichromate crystal are heated with conc. HCl
  - (a)  $O_2$  is evolved
  - (b) Chromyl chloride vapours are evolved
  - (c)  $Cl_2$  is evolved
  - (d) No reaction takes place
- Which of the following does not illustrate the 12. anomalous properties of lithium [MP PET 1993]
  - (a) The melting point and boiling point of Li are comparatively high
- (b) Li is much softer than the other group I metals
  - (c) Li forms a nitride  $Li_3N$  unlike group I metals
  - (d) The ion of Li and its compounds are more heavily hydrated than those of the rest of the group
- Correct order of increasing activity is
  - (a) Cu, Mg, Na
- (b) Na, Mg, Cu
- (c) Mg, Na, Cu
- (d) Cu, Na, Mg
- On heating anhydrous  $Na_2CO_3$ , ...... is evolved

[CPMT 1971, 79]

- (a) CO<sub>2</sub>
- (b) Water vapour
- (c) CO
- (d) No gas
- Chile saltpetre is

- [DPMT 1984; CPMT 1986, 89; CET Pune 1998; MP PMT 2003]
- (a) NaNO<sub>3</sub>
- (b)  $Na_2SO_4$
- (c)  $KNO_3$
- (d)  $Na_2SO_3$
- 16. A mixture of KCl and KF is added to sodium chloride
  - (a) To increase the conductivity of NaCl
  - (b) To decrease the melting point of NaCl
  - (c) To supress the degree of dissociation of NaCl
  - (d) To decrease the volatility of NaCl
- A well known reagent which contains copper sulphate, sodium potassium tarterate and sodium hydroxide is
  - (a) Fenton's reagent
- (b) Schiff's reagent
- (c) Fehling's solution
- (d) Nessler's reagent

Sodium metal can be stored under

[CPMT 1972, 85; BHU 1983]

- (a) Benzene
- (b) Kerosene
- (c) Alcohol
- (d) Toluene

19.		method of preparing	31.	The colour of hydrogen	is [IIT 1980]	
		ne action of HCl and[JIPMER:	2000]	(a) Black	(b) Yellow	
	(a) Al	(b) <i>K</i>		(c) Orange	(d) None of these	
	(c) Fe	(d) Zn	<b>32.</b>		wing salts gives aqueous	
20.		and other considerations		solution which is weakly		
	is expected to have the highest melting point[AIEEE 2005]			(a) NaHCO <sub>3</sub>	(b) NaHSO 4	
	(a) LiCl	(b) NaCl		(c) NaCl	(d) $NH_4HCO_3$	
	(c) KCl	(d) RbCl	33.	An example for a double	e salt is [KCET 2002]	
21.	The correct formula of h			(a) Silver nitrate	(b) Mohr's salt	
		(b) $Na_2SO_4$		•	ide (d)Cupromonium sulphate	
	(c) $Na_2S_2O_3.4H_2O$		34.	flame of Bunson burner		
22.	_	aly used to determine		(a) Low ionization poter	ntial	
		netrically is [AIIMS 2003]		(b) Low melting point		
	<ul><li>(a) Oxalic acid</li><li>(b) Disodium salt of ED?</li></ul>	T.A		(c) Softness		
	(c) Sodium citrate	IA	orbit	' '	lectron in the outermost	
	(d) Sodium thiosulphate			Which of the following i	s the smallest cation	
23.	$K_2CS_3$ can be called pot		33.	winen of the following i	[MP PMT 1993]	
_5.	(a) Thiocyanate	(b) Thiocarbonate		(a) <i>Na</i> +	(b) $Mg^{+2}$	
	(c) Thiocyanace	(d) Sulphocyanide		(c) $Ca^{+2}$	(d) $Al^{+3}$	
24.	Which is most basic in c		36.	` ,	may be arranged in the	
	(a) RbOH	(b) <i>KOH</i>	30.		their standard electrode	
	(c) NaOH	(d) LiOH		potentials as	[CPMT 1990]	
25.	When washing soda is h	eated [AFMC 2005]		(a) <i>K</i> , <i>Ca</i> , <i>Li</i>	(b) <i>Li</i> , <i>K</i> , <i>Ca</i>	
	<ul><li>(a) CO is released</li><li>(b) CO + CO<sub>2</sub> is released</li></ul>			(c) Li, Ca, K	(d) Ca, Li, K	
				Alkali metals lose electr	ons in [CBSE PMT 1990]	
	(c) $CO_2$ is released			(a) s-orbitals	(b) <i>p</i> -orbitals	
	(d) Water vapour is rele	eased		(c) d-orbitals	(d) <i>f</i> -orbitals	
26.	Which of the following i	s correct [CPMT 1971]	38.		acts with nitrogen directly	
	(a) All carbonates are so				ee 1992; MP PMT 2000; BHU 2000]	
	(b) Carbonates of Na, I	$K$ and $NH_4$ are soluble in		(a) Li	(b) <i>Na</i>	
wate				(c) K	(d) Rb	
		r, Ba are soluble in water	39.	water	has density greater than	
	(d) All carbonates are in				[MP PET 1994]	
27.	Nitre is	[CPMT 1986]		(a) Li	(b) <i>Na</i>	
	(a) $AgNO_3$	(b) $KNO_3$		(c) K	(d) <i>Rb</i>	
	(c) $NH_4NO_3$	(d) $NaNO_3$	40.	_	alkali metal sodium with	
28.	Nelson cell is used for the			water, is made use of	[MP PMT 1994]	
		[CPMT 1985]		(a) In drying of alcohols		
	(a) Slaked lime	(b) Baryta		(b) In drying of benzene		
	(c) Sodium	(d) Caustic soda		(c) In drying of ammoni		
29.	Potash alum is a	[CPMT 1986; MNR 1981]	41	(d) As a general drying a	_	
	<ul><li>(a) Complex salt</li><li>(c) Double salt</li></ul>	(b) Acid salt (d) Normal salt	41.	Which of the following h	ras smaller size [RPET 2003]	
30.		strial manufacturing of		(a) <i>H</i>	(b) He <sup>+</sup>	
Ju.	sodium carbonate is kno	own as [CPMT 1978, 86; MP PM	T 1995	$\begin{bmatrix} a & H \\ C & H^2 \end{bmatrix}$	(d) Li <sup>2+</sup>	
	(a) Castner process	(b) Haber's process		$(c)_{1}\pi$	(u) Li	
	(c) Le-blanc process	(d) Chamber process				

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42.		$HF$ to form $KHF_2$ . The		(a) LiCl has higher m	0 1	
	compound contains the	<del>-</del>		(b) LiCl dissolves in v		
	(a) $K^+$ , $F^-$ and $H^+$	(b) $K^+$ , $F^-$ and $HF$		(c) LiCl would ionize		
	(c) $K^+$ and $[HF_2]^-$			(d) Fused <i>LiCl</i> woul fused <i>NaCl</i>	d be less con	ducting than
43.	Which alkali metal is n	nost metallic in character [MH CET 2001]	52.	In the Castner's prosodium, the anode is		
	(a) <i>K</i>	(b) <i>Cs</i>		(a) Copper	(b) Iron	
	(c) Na	(d) <i>Li</i>		(c) Sodium	(d) Nickel	
44.	The property of hydrofrom other alkali metal	gen which distinguishes it is [MP PET 1996]	53.	Which of the followinitride	ing <i>s</i> -block el	ements forms
	(a) Its electropositive of	character				[RPET 2003]
	(b) Its affinity for non-	metals		(a) Ba	(b) <i>Be</i>	
	(c) Its reducing charac	ter		(c) <i>Ca</i>	(d) <i>Li</i>	
	(d) Its non-metallic cha	aracter	54.	Tincal is		Pb. PMT 2001]
45.	Which of the following high rate	ng reacts with water with		(a) $Na_2CO_3.10H_2O$	(b) $NaNO_3$	
		[AFMC 1995]		(c) NaCl	(d) $Na_2B_4O$	$_{7}.10H_{2}O$
	(a) Li	(b) <i>K</i>	55.	Which has minimum	solubility	[BHU 2003]
	(c) Na	(d) <i>Rb</i>		(a) $Br_2S_3$	(b) $Ag_2S$	
46.	The valence shell e	lectronic configuration of		(c) CoS	(d) PbS	
	alkali metals is	[MP PET 1996; UPSEAT 2001]	56.	Cryolite helps in		[BHU 2003]
	(a) $ns^2np^1$	(b) ns <sup>1</sup>		(a) Lowering the mel	~ -	
		• •		<ul><li>(b) Increasing the me</li><li>(c) Increasing the ele</li></ul>	~ -	
	(c) $(n-1)p^6 ns^2$	(d) $(n-1)d^2ns^2$		(d) Decreasing the ele		
47.	Alkali metals are	[MP PMT 1996]	57.	In certain matters		-
	(a) Li, Na, Be, Mg, Cs	(b) <i>Li, Na, K, Rb, Cs</i>	3/•	alkali metals, the mai		
	(c) Na, K, Mg, Ca, Rb	(d) <i>K, Rb, Cs, Ba, Sr</i>		(a) Small size of <i>Li</i> at		
48.		an element is 11. Its oxide		(b) Extremely high el		
	will be			(c) Greater hardness	of Li	
		[MP PMT 1996]		(d) Hydration of $Li^+$	ion	
	(a) Acidic	(b) Basic	58.	Acidified potassium		solution is
	(c) Acid and basic both			decolourised by		
49.	-	action of sodium carbonate		(a) Bleaching powder	(b) Microco	smic salt
	is done by	[CPMT 1982; MP PMT 1996]		(c) Mohr salt	(d) White v	
	(a) Lead-chamber proc		59.	Which one of the disinfectant in water	_	
	(b) Haber's process	C35		(a) Alum	(b) Charcoa	[NDA 1999]
	(c) Solvay's process			(c) Kieselguhr	(d) Potassii	
	(d) Castner's process		pern	nanganate	(4) 1044551	****
50.	Alkali metals are stron	g reducing because	60.	Sodium thiosulphate	is used in photo	graphy
50.	(a) These are monovale					[UPSEAT 1999]
	(b) Their ionisation po			(a) To convert metall	ic silver into sil	ver salt
	-	• •		(b) AgBr grain is redu	iced to non-met	allic silver
	much negative	ectrode potential are very		(c) To remove reduce		
	(d) These are metals			(d) To remove undecomplete $Na_3[Ag[S_2O_3)_2]$ (a		n the form of
51.		g statement about <i>LiCl</i> and	<i>c</i> -			CDATE DOOL 5 13
	NaCl is correct	[Kurukshetra CEE 2002]	61.	Composition of borax (a) $Na_2B_4O_7.4H_2O$	(b) $Na_2B_4O_2$	<b>SEAT 2001;04]</b> <sub>7</sub> .10 <i>H</i> <sub>2</sub> <i>O</i>

#### 758 s and p-Block Elements (c) $NaBO_{2}$ (d) $Na_2BO_3$ (b) Does not show regular variation (c) Increases as we go down the group 62. When sodium dicarbonate is heated strongly for calcined in a kiln, it forms[CPMT 2000; KCET (Med.) 2000] (d) Decreases as we go down the group (b) $Na_2CO_3$ Which of the following properties is not true for an alkali metal [Pune CET 1998] (c) NaCO<sub>3</sub> (d) NaHCO 3 (a) Low atomic volume The strongest reducing agent is [MP PET 2001] (b) Low ionization energy (a) K (b) Al (c) Low density (c) Mq (d) Br (d) Low electronegativity The word 'alkali' is used for alkali metals 64. indicates Which of the following alkali metals has the 73. [RPMT 1999] biggest tendency for the half reaction, [DPMT 2001] (a) Ash of the plants (b) Metallic nature $M(g) \rightarrow M^+(aq) + e^-$ (c) Silvery lusture (d) Active metal (a) Lithium (b) Sodium 65. Potassium nitrate is called [RPMT 1999] (d) Potassium (c) Cesium (a) Mohr's salt (b) Gypsum Which one of the following metalic hydroxides (c) Indian salt petre (d) Chile salt petre does not dissolve in sodium hydroxide solution [KCET (Me Which of the following chemicals, in addition to (b) $Al(OH)_2$ (a) $Zn(OH)_2$ water, are used for the manufacture of $Na_2CO_3$ by (c) $Fe(OH)_3$ (d) $Pb(OH)_2$ Solvay process Which one of the following on heating will not [Roorkee 1999] give $CO_2$ (a) NaCl, CO and $NH_3$ [NDA 1999; BHU 2000] (b) $NaCl, CO_2$ and $NH_3$ (a) CaCO<sub>3</sub> (b) $Na_2CO_3$ (c) NaCl, $NH_4Cl$ and $CO_2$ (c) $PbCO_3$ (d) $Li_2CO_3$ (d) $NaHCO_3$ , CO and $NH_3$ **76.** *NaOH* is prepared by the method [AFMC 2005] Which metal forms amide with $NH_3$ at $300^{\circ} C$ (a) Down's cell (b) Castner cell [CPMT 1994] (c) Solvay process (d) Castner Kellner cell (a) Mg (b) Pb Sodium gives blue colour with $NH_3$ solution, this (c) Al (d) Na blue colour is due to 68. When sodium is heated with moist air, then the [UPSEAT 2000,02; AMU 2002; RPMT 2002] product obtained is [AIIMS 1999] (a) Ammoniated $Na^{\oplus}$ (b) Ammoniated $Na^{\Theta}$ (a) $Na_2O$ (b) NaOH (c) Ammoniated $e^-$ (d) $Na^+/Na^-$ pair (c) $Na_2CO_3$ (d) $Na_2O_2$ The strongest reducing agent of the alkali metal is [CPMT 1999; Pb.CET 2001] inorganic compound first 69. melts then (b) Na resolidifies and then liberates a gas. It may be[DPMT 2002] (a) Li (c) K (d) Cs (b) $Al_2O_3$ (a) $MnO_2$ With the increase in atomic weights, melting (c) $KMnO_4$ (d) KClO<sub>3</sub> points of the alkali metals [MP PMT 1995] On dissolving moderate amount of sodium metal (a) Increase in liquid $NH_3$ at low temperature, which one of (b) Decrease the following does not occur [AIIMS 2003] (c) Remain constant (a) Blue coloured solution is obtained (d) Do not show definite trend 80. The reaction of water with sodium and potassium (b) $Na^+$ ions are formed in the solution (c) Liquid $NH_2$ becomes good conductor of [BHU 1999] electricity (a) Exothermic (d) Liquid ammonia remains diamagnetic (b) Endothermic The solubility of the alkali metal carbonates 71. (c) Reversible

[Pune CET 1998]

(a) Increases at first and then decreases

(d) Irreversible and endothermic

81.	_	cyanide crystals are heated		(c) $CH_3COONa$ (d) None of these
	with concentrated sulphuric acid, the gas evolved is		91.	A fire of lithium, sodium and potassium can be extinguished by [DCE 2003]
		E PMT PMT 1999; KCET 2000]		(a) $H_2O$ (b) Nitrogen
	(a) Ammonia	(b) Sulphur dioxide		(c) $CO_2$ (d) Asbestos blanket
0-	(c) Carbon dioxide	(d) Carbon monoxide		<del>-</del>
82.	Characteristic feature	of alkali metals is  [RPMT 2000; MP PMT 2004]	92.	Which of the following metal has stable carbonates
	(a) Good conductor of heat and electricity			[AFMC 2004]
	(b) High melting point	s		(a) Na (b) Mg
	(c) Low oxidation pote	entials		(c) Al (d) Si
	(d) High ionization pot	tentials	93.	Aluminium reacts with caustic soda to form [DCE 20
83.		ompound of an element of		(a) Aluminium hydroxide
	group IA the substance $X$ gives a violet colour in			(b) Aluminium oxide
	flame test, X is			(c) Sodium meta-aluminate
		35, 86; CPMT 1985; DCE 2000]		(d) Sodium tetra aluminate
	(a) LiCl	(b) NaCl	94.	Alkaline earth metals are denser than alkali metals,
Q A	(c) KCl Which of the follow	(d) None		because metallic bonding in alkaline earth's metal,
84.	Which of the following alkali metal ions has lowest ionic mobility in aqueous solution [KCET 2000]			is [BHU 2004]
	(a) $Rb^+$ (b) $Cs^+$		•	(a) Stronger (b) Weaker
	(c) $Li^+$	(d) Na <sup>+</sup>		(c) Volatile (d) Not present
Q =		rities to magnesium in its	95.	Which of the following is a false statement [CPMT 2
85.	chemical behaviour be	•		(a) Fluorine is more electronegative than chlorine
		ater electronegativity and		(b) Nitrogen has greater IE <sub>1</sub> than oxygen
	similar polarizing power.  (b) Similar size same electronegativity and lower polarizing power  (c) Similar size, same electronegativity and			(c) Lithium is amphoteric
				(d) Chlorine is an oxidising agent
			96.	Which is most basic in character [UPSEAT 2004]
				(a) CsOH (b) KOH
	similar high polari	zing power		(c) NaOH (d) LiOH
	(d) None of these		97.	Photoelectric effect is maximum in [AFMC 2004]
86.		following is the most		(a) Cs (b) Na
	electropositive elemen			(c) <i>K</i> (d) <i>Li</i>
	(a) Calcium	(b) Chlorine	98.	A metal $M$ reacts with $N_2$ to give a compound 'A'
o-	(c) Potassium	(d) Carbon		$(M_3N)$ . 'A' on heating at high temperature gives
87.	the formation of	sodium chloride leads to [KCET 1990]		back 'M' and 'A' on reacting with $H_2O$ gives a gas
	(a) $Na$ and $H_2$	(b) $Na$ and $O_2$		$^{\prime}B^{\prime}.$ $^{\prime}B^{\prime}$ turns $^{\prime}CuSO_4$ solution blue on passing
	_	(d) $Na$ and $Cl_2$		through it. A and B can be [DCE 2003]
88.		nate is heated the product		(a) $Al$ and $NH_3$ (b) $Li$ and $NH_3$
00.	obtained is	nate is heated the product		(c) $Na$ and $NH_3$ (d) $Mg$ and $NH_3$
		[Pb. CET 2000; DCE 2004]	99.	A solid compound $X'$ on heating gives $CO_2$ gas
	(a) Na	(b) $Na_2CO_3$		and a residue. The residue mixed with water
	(c) NaCO <sub>3</sub>	(d) $Na_2(HCO_3)$		forms 'Y'. On passing an excess of ${\it CO}_2$ through 'Y'
0،				in water, a clear solution, ${}^{\prime}Z^{\prime}$ is obtained. On
89.	_	(b) Pleashing slethes		boiling $'Z'$ , compound $'X'$ is reformed. The
	(a) Making explosives (b) Bleaching clothes			compound 'X' is [CBSE PMT 2004]
	(c) Water softening	(d) All of the above		(a) $Na_2CO_3$ (b) $K_2CO_3$
00	Which of the following salt does not get			(c) $Ca(HCO_3)_2$ (d) $CaCO_3$
90.	hydrolysed in water			
90.	hydrolysed in water	[CPMT 2004]	100.	mongst $LiCl$ , $RbCl$ , $BeCl$ <sub>2</sub> and $MgCl$ <sub>2</sub> the compounds

		[Pb. CET 2004]	111.	When NaCl is dissolve	ed in water, the sodium ion	
	(a) LiCl and RbCl	(b) $MgCl_2$ and $BeCl_2$		is		
	(c) $RbCl$ and $BeCl_2$	(d) $RbCl$ and $MgCl_2$			[CPMT 1989]	
101.	Salt cake is			(a) Oxidised	(b) Reduced	
	(a) Sodium sulphate			(c) Hydrolysed	(d) Hydrated	
	(b) Sodium chloride			Sodium metal cannot b	e stored under	
	(c) Sodium bisulphite				[CPMT 1985, 88, 94]	
	(d) Sodium sulphate and	l Sodium chloride		(a) Benzene	(b) Kerosene	
102.	Globar salt is	[BHU 1983; CPMT 1988, 91;		(c) Alcohol	(d) Toluene	
	(a) M (ii) 7H (ii)	IIT 1985; MP PET 2000]	113.	-	is used for the preparation	
	(a) $MgSO_4.7H_2O$	(b) $CuSO_4.5H_2O$		of	[CDMT 1095, DIIII 1096]	
	(c) $FeSO_4.7H_2O$	(d) $Na_2SO_4.10H_2O$		(a) Caustic soda	[CPMT 1985; BHU 1986] (b) Caustic potash	
103.	The colour given to the f	-			(d) Slaked lime	
	(a) Light red	[CPMT 1980; MP PET 1986] (b) Golden yellow	11.4	(c) Baryta		
	(c) Green	(d) Pink	114.	$200^{\circ}C$ , it forms	over solid <i>NaOH</i> heated to [MP PMT 1985]	
104.	Solvay's process is used			(a) $Na_2CO_3$	(b) NaHCO <sub>3</sub>	
	corrug o process is used	[CPMT 1982; AIIMS 1987]			3	
	(a) Ammonia	(b) Sodium bicarbonate		(c) HCOONa	(d) None	
	(c) Sodium carbonate	(d) Calcium carbonate	115.		ed by electrolysis of brine of the reaction are[KCET 1990]	
105.	Sodium when heated in	a current of dry ammonia		(a) $Cl_2$ and $H_2$	(b) $Cl_2$ and $Na - Hg$	
	gives				_	
	(a) Sodium nitrite	[NCERT 1981; KCET 2000]		(c) $Cl_2$ and $Na$	(d) $Cl_2$ and $O_2$	
	(c) Sodium amide	<ul><li>(b) Sodium hydride</li><li>(d) Sodium azide</li></ul>	116.		manufactured by Solvay	
106.	Washing soda is	(d) Sourum azide			hat are recycled are[KCET 1993; DC	
	•	OPMT 1982; CBSE PMT 1990;		(a) $CO_2$ and $NH_3$	(b) $CO_2$ and $NH_4Cl$	
	,	MP PMT 1987, 96]		(c) NaCl, CaO	(d) $CaCl_2$ , $CaO$	
	(a) $Na_2CO_3.10H_2O$	(b) $Na_2CO_3.H_2O$	117.		cts, obtained in the Solvay	
	(c) $Na_2CO_3.5H_2O$	(d) $Na_2CO_3$			ng sodium carbonate, are[KCET 198	
107.	The substance used to d	ecolourise and purify oils		(a) Quick lime and $CO_2$		
	is	•		(b) $NaHCO_3$ and $NH_4C$	T .	
		[MP PMT 1987]		(c) $NH_4Cl$ solution and	d quick lime	
	(a) Sodium carbonate	(b) Sodium chloride		(d) $NaHCO_3$ and $CO_2$		
0	(c) Sodium hydroxide	(d) Sodium sulphate	110		andium carbonate which of	
108.		sea water is[MP PMT 1998]	110.	the following is used	sodium carbonate, which of [AFMC 1992]	
	(a) $MgCl_2$	(b) NaCl		(a) Slaked lime	(b) Quick lime	
	(c) $MgSO_4$	(d) $CaSO_4$		(c) Lime stone	(d) NaOH	
109.		exhibited by sodium is	119.		are left in open air, they	
	explained by	[IIT 400=1			ound each crystal as[CPMT 1974]	
	(a) Diffusion of sodium	[IIT 1987]		(a) They start melting		
	(b) Oscillation of loose 6			(b) They absorb moistu	ıre from air	
	(c) Excitation of free pr			(c) They react with air	to form a liquid compound	
	(d) Existence of body ce			(d) They absorb $CO_2$ fr	rom air	
110.		ets with water at room	120.	Sodium carbonate rea	cts with $SO_2$ in aqueous	
	temperature is			medium to give	2 1	
	=	MP PMT 1996; MP PET 1998]		0 -	[MP PMT 1982, 85]	
	(a) Copper	(b) Iron		(a) NaHSO <sub>3</sub>	(b) $Na_2SO_3$	
	(c) Magnesium	(d) Sodium		(c) NaHSO 4	(d) $Na_2SO_4$	
				(5) 1101150 4	(5) 114250 4	

121.	Baking soda is	[CPMT 1974, 78, 79, 91;		(b) 10% solution of	КОН	
	BHU 1979; M	anipal MEE 1995; AIIMS 1996;		(c) 10% solution of	$Ca(OH)_2$	
	CPMT 1973; RPET 1999;	AFMC 2001, 05; Pb. CET 2002]		(d) 10% solution of	$Na_{2}CO_{3}$	
	(a) $Na_2CO_3$	(b) NaHCO <sub>3</sub>	131.		w colour to Bunsen flame	
	(c) $Na_2SO_4$	(d) $K_2CO_3$	131.	because of	w colour to Builsen Hame	
122.	Soda ash is	[KCET 1993]			[RPMT 1997]	
	(a) $Na_2CO_3.H_2O$	(b) NaOH		(a) Low ionisation po	otential	
	(c) $Na_2CO_3$	(d) NaHCO <sub>3</sub>		(b) Sensitivity		
122.	Soda lime is	[KCET 1993]		<ul><li>(c) Sublimation</li><li>(d) Absorbed high ra</li></ul>	diation	
5	(a) NaOH	(b) <i>CaO</i>	132.	•	excess of <i>NaOH</i> solution, the	
	(c) NaOH and CaO	(d) $Na_2CO_3$	13=1	compound obtained i		
124		in nuclear reactors to[KCET 19	<b>Q</b>	(a) $Sn(OH)_2$	(b) $Na_2SnO_3$	
124.		n order to control the chain	09]	(c) $Na_2SnO_2$	(d) $SnO_2$	
react		r order to control the chain	133.	Identify the correct s	statement [CPMT 1997]	
	(b) Slow down the fast	neutrons		•	ım can be prepared and	
	(c) Absorb the heat generated by nuclear fission				rolysing an aqueous solution	
	(d) Extract radio-isoto	pes produced in the reactor		of sodium chlorid		
125.	Squashes are stored by	adding [AFMC 1989]			n is a strong oxidizing agent n is insoluble in ammonia	
	(a) Citric acid	(b) <i>KCl</i>		(d) Elemental sodium		
	(c) $Na_2SO_3$	(d) Sodium	13/1.	Calcium is obtained b	·	
meta	bisulphite		-5-7-1	(a) Roasting of limes	•	
126.	Sodium thiosulphate	$(Na_2S_2O_3.5H_2O)$ is used in		•	olution of calcium chloride in	
	photography to	[CPMT 1972, 74, 79;		$H_2O$		
	DPMT 198	3; Bihar CEE 1995; MNR 1995]		(c) Reduction of calc	ium chloride with carbon	
	(a) Reduce silver brom	nide to metallic silver		_	molten anhydrous calcium	
	(b) Convert metallic si	lver to silver salt	chlo			
		posed $AgBr$ as a soluble	135.		ride solution is electrolysed, ted at the cathode is[Kurukshetra (	
	silver thiosulphate	_		(a) Oxygen	(b) Hydrogen	
	(d) Remove unreduced			(c) Chlorine	(d) Air	
127.	Which of the following	pair can't exist in solution	136.		sis of fused sodium chloride,	
	(a) N-HCO and N-OI	[IIT 1986; DCE 1999]		the anodic reaction is [KCET 1998]		
	-	$H$ (b) $Na_2CO_3$ and $NaOH$		(a) Reduction of sodi		
	(c) $Na_2CO_3$ and $NaCl$	(d) $NaHCO_3$ and $NaCl$		(b) Oxidation of sodi		
128.	Sodium thiosulphate is	s prepared by [IIT 1996]		(c) Reduction of chlo		
	(a) Reducing $Na_2SO_4$ s	solution with $H_2S$	125	(d) Oxidation of chlo	ng does not participate in the	
	(b) Boiling $Na_2SO_3$ so	olution with $S$ in alkaline	13/.		the manufacture of $Na_2CO_3$ [EAMC]	
medi	um			(a) $NH_3$	(b) <i>NaCl</i> solution	
	(c) Neutralising $H_2S_2$	$O_3$ solution with $NaOH$		(c) $CO_2$	(d) $H_2SO_4$	
	(d) Boiling $Na_2SO_3$ s	solution with $S$ in acidic	128	_	recipitate produced by adding	
	um		130.	NaOH solution to $Hg$		
medi			061			
medi <b>129.</b>	When NaOH is prepar	ed, the gas released is[CPMT 19	190]	(a) renow	(b) Black	
	When $NaOH$ is prepartable (a) $Cl_2$	ed, the gas released is [CPMT 19 (b) $H_2$	190]	<ul><li>(a) Yellow</li><li>(c) Brown</li></ul>	(b) Black (d) White	
	(a) Cl <sub>2</sub>	(b) H <sub>2</sub>		(c) Brown		
129.		_		(c) Brown The cell used for the	(d) White	

#### 762 s and p-Block Elements (c) Solvay cell (d) Nelson cell To remove last traces of water from alcohol, the metal used is **140.** Slaked lime $[Ca(OH)_2]$ is used in the manufacture (a) Sodium (b) Potassium [UPSEAT 2000] (c) Calcium (d) Aluminium (a) Cement (b) Fire bricks Plaster of Paris is (c) Pigment (d) Medicine 2. [CPMT 1972, 76, 78, 83, 87, 88, 90, 91, 93, 94; JIPMER 2002; 141. The alum used for purifying water is [KCET (Med.) 2001] MP PET 1986, 2001; BHU 1992, 95, 2000; MNR 1982; DCE 2000; (a) Ferric alum (b) Chrome alum Manipal MEE 1995; NCERT 1976; Bihar MEE 1997; EAMCET 1978; (d) Ammonium alum (c) Potash alum AMU 1982, 84; DPMT 1982, 83] 142. Which one of the following metalic hydroxides does not dissolve in sodium hydroxide solution [KCET (Med.) 2001] (a) $CaSO_4.2H_2O$ (b) $CaSO_4.3H_2O$ (d) $CaSO_4 \cdot \frac{1}{2}H_2O$ (b) $Al(OH)_3$ (a) $Zn(OH)_2$ (c) $CaSO_A.H_2O$ (c) $Fe(OH)_2$ (d) $Pb(OH)_2$ Which of the following substance is used as 3. 143. In which of the following processes, fused sodium dehydrating agent in laboratory [MP PMT 1987] hydroxide is electrolysed at a 330°C temperature (a) Calcium chloride (b) Sodium chloride for extraction of sodium [CBSE PMT 2000; AFMC 2001] (c) Sodium carbonate (d) Potassium nitrate (a) Castner's process (b) Down's process The metal that is extracted from sea water is 4. [EAMCET 1978; CPMT 1988; CET Pune 1998; (c) Cyanide process (d) Both (b) and (c) MP PET 2000] **144.** Excess of $Na^+$ ions in our system causes (a) *Ba* (b) Mg [KCET (Med.) 2001] (c) *Ca* (d) Sr (a) High B.P. (b) Low B.P. 5. Which of the following ore contains both (c) Diabetes (d) Anaemia magnesium and calcium [MDAT Bihar 1984; MP PET 2003] **145.** Ferric composition alum the (b) Dolomite (a) Magnesite [Orissa JEE 2002] $(NH_4)_2SO_4.Fe_2(SO_4)_3.xH_2O$ (c) Carnellite (d) Phosphorite (a) 7 (b) 24 6. Epsom salt is (c) 6(d) 15 [EAMCET 1978, 80; BHU 1979; MP PET 1999; **146.** If *Na* is heated in presence of air, it forms [AFMC 2002] CPMT 1988, 89, 90; Bihar MEE 1996] (a) $CaSO_4.2H_2O$ (b) $BaSO_4.2H_2O$ (a) $Na_2CO_3$ (b) $Na_2O_2$ (d) Both (b) and (c) (c) $Na_2O$ (c) $MgSO_4.2H_2O$ (d) $MgSO_4.7H_2O$ 147. Which of the following is most reducing agent[RPMT 2502] Setting of plaster of paris is[MP PMT 1985; CPMT 1989] (a) Oxidation with atmospheric oxygen (a) $HNO_3$ (b) Na (b) Combination with atmospheric $CO_2$ (d) Cr (c) Cl<sub>2</sub> (c) Dehydration 148. Pyrolusite is [DPMT 2002] (d) Hydration to yield another hydrate (a) Carbonate ore (b) Sulphur ore prevent magnesium from oxidation in 8. (c) Silicon ore (d) None of these electrolytic extraction process 149. In the manufacture of metallic sodium by the (a) Some calcium fluoride is added fused salt electrolysis (Down's process) a small amount of calcium chloride is added to[мр рет 1993; мр рмт робрате chlorides are added (a) Improve the electrical conduction (c) Metal is taken out by spoons (d) The whole process is done in an atmosphere (b) Increase the temperature of electrolysis of coal gas (c) Bring down the melt temperature Which of the following metal is found in green (d) Stabilize the metallic sodium colouring pigment chlorophyll of plants **150.** Sodium metal is extracted by [MP PMT 1996] [KCET 1993; RPMT 1999; MP PET 2002] (a) Electrolysis of aqueous solution of sodium (a) *Fe* (b) Mg chloride (c) Na (d) Al (b) Electrolysis of fused sodium chloride Which of the following metal carbonate is (c) Heating sodium oxide with carbon decomposed on heating [MNR 1985; MP PET 1994; Pb. CET 2 (d) Heating sodium oxide with hydrogen (a) $MgCO_3$ (b) $Na_2CO_3$

(c)  $K_2CO_3$ 

(d)  $Rb_2CO_3$ 

Alkaline earth metals

11.		ic configuration of alkaline				[NCERT 1982]
	earth metal is				f beryllium carbide	
		o; CPMT 1985, 93; MP PAT 1993]		(b) Electrolysis of	f fused beryllium cl	nloride
	(a) ns <sup>2</sup>	(b) ns <sup>1</sup>		(c) Reduction of b	peryllium oxide wit	h carbon
	(c) $np^{6}$	(d) $nd^{10}$			of beryllium	halide with
12.	Metallic magnesium	is prepared by	mag	<b>BA 6</b> <sup>i</sup> 1973, 77]		
	(a) Reduction of $MgO$ by coke			Mark the incorrec	t statement	
	(b) Electrolysis of ac	queous solution of $Mg(NO_3)$		=	cheap and possess	good covering
	(c) Displacement of	$f$ $Mg$ by iron from $MgSO_4$	pow			
solu	_			(b) Lithopone is y		
	(d) Electrolysis of m	olten MgCl <sub>2</sub>		(c) Lithopone is sulphide and a	prepared by m	ixing barium
13.	Of the metals $Be$ , $A$	Mg, $Ca$ and $Sr$ of group II A.		-	-	aulphata and
•		the least ionic chloride would		zinc sulphide	n mixture of barium	i suipiiate and
	be formed by		22.	_	$MgCl_2$ can be prepared	ared from the
		[NCERT 1980; CPMT 1980]		hydrated salt by	=	MP PMT 1989]
	(a) <i>Be</i>	(b) <i>Mg</i>		(a) Heating the hy		MI 1MI 1909]
	(c) <i>Ca</i>	(d) <i>Sr</i>			ydrate with $M_g$ rib	hon
14.	Which one of the following is fluorspar				_	)DOII
	(a) $CaF_2$ (b) $CaO$			(c) Melting the hy		
	(c) $H_2F_2$	(d) CaCO <sub>3</sub>		atmosphere of	hydrate to red f <i>HCl</i> gas	neat in an
15.	Which one is known	as barytes [CPMT 1987]	23.	-	is obtained by the	interaction of
	(a) $BaSO_4$	(b) $BaCl_2.2H_2O$			MT 1972, 78, 89; 200	
	(c) <i>BaO</i>	(d) <i>BaCO</i> <sub>3</sub>		(a) Conc. solution	of $Ca(OH)_2$	
16.	Which of the followi	ng sulphates have the highest		(b) Dilute solution	n of $Ca(OH)_2$	
10.	solubility in water[EAMCET 1980,84,85; MP PMT 1994;			(c) Dry calcium of	xide	
	Kurukshetra CEE	1998; AFMC 1990; MP PET 1994]		(d) Dry slaked lim	ne	
	(a) $MgSO_4$	(b) $BaSO_4$	24.	Deep pink colour	is given to flame by	the salts of
	(c) CaSO <sub>4</sub>	(d) BeSO $_4$		(a) Strontium	(b) Potassii	um
17.	The composition for	mulae of gypsum is		(c) Zinc	(d) Barium	
-/•	-	75, 78, 82; DPMT 1982; IIT 1978;	25.	_	ve which colour w	hen put in a
		1981; MP PMT 1996; RPMT 1997]		flame		
	(a) $(CaSO_4)_2.H_2O$	(b) 2 <i>CaSO</i> <sub>4</sub>		(a) Brick red	(b) Green	
	(c) $CaSO_4.2H_2O$	(d) $2CaSO_4.H_2O$		(c) White	(d) Pink	•
10		of [EAMCET 1998; AIIMS 2000]	26.	Phosphine is obta		_
18.				(a) Calcium suner	rphosphite (b)Calci	[Roorkee 1995] ium phosphide
	(a) $CaCO_3$ , sand and			-	osphide (d) Calcium	
	(b) Slaked lime and			hypophosphid		•
	(c) Slaked lime, sand and water			Calcium is obtained by		
	(d) $CaCO_3$ and $CaO$			[DPMT 1980	; IIT 1980; CPMT 199	6;AIIMS 2001]
19.	Gypsum CaSO <sub>4</sub> .2H <sub>2</sub> O	O on heating to about $120^{\circ} C$		(a) Roasting of lir	ne stone	
	forms a compound	d which has the chemical		(b) Reduction of	CaCl <sub>2</sub> with carbon	
	composition represe	nted by[CPMT 1978, 82, 88, 90; E	AMCET	1978; Electrolysis of	f a solution of <i>CaCl</i>	<sub>2</sub> in water
		DPMT 1982, 83; NCERT 1979]		(d) Electrolysis of		
	(a) $CaSO_4$	(b) $2CaSO_4.H_2O$	28.	_	ossesses biggest ato	mic radii
	(c) $CaSO_4.H_2O$	(d) $2CaSO_4.3H_2O$	20.	(a) P	(b) Si	inic raum
20.	The highly effici	ent method of obtaining		(c) Al	(d) Mg	
	beryllium is		29.	Magnesia is	() 6	
			- 5.			

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	(a) MgCO <sub>3</sub>	(b) <i>MgO</i>		(a) Aluminium	(b) Boron
	(c) $MgSO_4$	(d) $MgCl_2$		(c) Magnesium	(d) Sodium
30.		l <i>MgO</i> is called [ <b>DPMT 1984</b> ]	41.	The right order of the alkaline earth metals i	e solubility of sulphates of n water is
	(a) Double salt	(b) Sorrel cement		[MP PET 199	93; Pb. CET 2000; DPMT 2004]
	(c) Portland cement			(a) $Be > Ca > Mg > Ba >$	Sr
31.	-	[AFMC 1992; BHU 1983, 86, 95;		(b) $Mg > Be > Ba > Ca >$	Sr
	(a) $BaO + ZnSO_4$	JIPMER 1999; RPET/PMT 1999] (b) $ZnO + BaSO_4$		(c) $Be > Mg > Ca > Sr >$	Ba
		(d) $ZnS + BaSO_4$		(d) $Mg > Ca > Ba > Be >$	Sr
	•	·	42.	Which of the follow	ring has highest electrode
32.	For bleaching powder	[EAMCET 1984; CPMT 1985]	-	potential	
	(a) Reacts with dilute	e acid to release chlorine			[CPMT 1990]
	(b) Oxidising agent	s deru to resease emerme		(a) <i>Be</i>	(b) <i>Mg</i>
	(c) Light yellow color	ıred powder		(c) <i>Ca</i>	(d) <i>Ba</i>
	(d) Highly soluble in	-	43.		etals $Ba$ , $Sr$ , $Ca$ and $Mg$
33.	Bleaching powder i molecular formula	s a compound having the		first ionisation potenti	
	[CPMT 1986, 89, 90	), 93; MP PMT 1996; BHU 2005]		(a) <i>Mg</i> , <i>Ca</i> , <i>Sr</i> , <i>Ba</i> ,	(b) Ca, Sr, Ba, Mg
	(a) CaOCl <sub>3</sub>	(b) CaOCl <sub>2</sub>		(c) Sr, Ba, Mg, Ca	(d) Ba, Mg, Ca, Sr,
	(c) CaClO	(d) CaClO <sub>3</sub>	44.		ring alkaline earth metals
34.	Calcium cynamide is	[CPMT 1986, 93]			s similar to aluminium[BHU 1983]
	(a) CaCHNH <sub>2</sub>	(b) CaCN <sub>2</sub>		(a) <i>Be</i>	(b) <i>Ca</i>
	(c) $CaC_2N_2$	(d) $Ca(CN)_2$	4-	(c) Sr	(d) Ba
35.	Which one of the follo	owing is a true peroxide	45.	hydroxide in water	g ions forms highly soluble [CPMT 1974, 76, 79, 82]
		999; CPMT 1981; Roorkee 1995]		(a) K <sup>+</sup>	(b) $Zn^{++}$
	(a) $SO_2$	(b) <i>BaO</i> <sub>2</sub>		(c) $Al^{+++}$	(d) Ca ++
	(c) $MnO_2$	(d) $NO_2$	46.	* *	soluble in water whereas
36.		ing is not a water absorber tance[CBSE 1989; JIPMER 2002]	-	barium sulphate is spa	aringly soluble because [IITJEE 1989] argy of $Na_2SO_4$ is less than
	(a) Silica gel	(b) $P_2O_5$		its lattice energy	
	(c) Conc. $H_2SO_4$	(d) Aqueous $CaCl_2$		(b) The hydration ener	rgy of $Na_2SO_4$ is more than
37•	The dark red colour to the presence of	of bombs in fireworks is due [Roorkee 1989; DPMT 2001		its lattice energy (c) The lattice energy	of $BaSO_4$ is more than its
		985; AFMC 1989; Roorkee 1989]		hydration energy	
	(a) <i>Na</i>	(b) <i>Ba</i>		_	y has no role to play in
	(c) Sr	(d) <i>K</i>	47.	solubility Which one of the follow	wing is most basic [CPMT 1977, 83]
38.	The most electropose earth metals is	sitive amongst the alkaline	4/•	(a) $Al_2O_3$	(b) <i>MgO</i>
	(-) D11'	[MP PMT 1993]		(c) $SiO_2$	(d) $P_2O_5$
	<ul><li>(a) Beryllium</li><li>(c) Calcium</li></ul>	(b) Magnesium (d) Barium	48.	_	are light and strong and so acture of aeroplane parts[EAMCET :
39.		ng salts is insoluble in water	DMT 4	(a) <i>Cr</i>	(b) <i>Sn</i>
	(a) CaCl <sub>2</sub>	but soluble in boiling water[MP (b) BaCl,		• •	(d) Mg
	(c) SrCl <sub>2</sub>	(d) $PbCl_2$	49.	works used give gree	sion of marriages, the fire n flame. Which one of the be present[CPMT 1980; AFMC 1989
40.		beryllium is approximately		10110 Willia Laureuro illay	MP PET 2002]
	equal to that of	Fren masm		(a) <i>Na</i>	(b) <i>K</i>
		[MP PMT 1993]			

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	(c) <i>Ba</i>	(d) <i>Ca</i>			(a) Li, Be, K, Mg, Ca	(b) Be, Mg, Ca, Sr, Ba
50.	$CaCO_3 \square CaO + CO_2$ reaction in a line goes to completion because [AFMC 2005]			61.		(d) Be, Mg, Ca, K, Rb ving substances is used in the lrying of neutral gases[AIIMS 1998]
	(a) CaO does not react to CO <sub>2</sub> to give CaCO <sub>3</sub>				(a) Sodium phospha	
	(b) Backward reaction	•			(b) Phosphorus pen	
	(c) $CO_2$ formed escape	es out			(c) Sodium sulphate	2
	(d) None of these				(d) Anhydrous calci	um chloride
51.	The wire of flash bulb		[CPMT 1988]	62.		ring can be represented by the
	(a) <i>Mg</i>	(b) <i>Cu</i>			configuration [Kr] 5 s	<sup>2</sup> ? [MP PMT 1997]
	(c) <i>Ba</i>	(d) Ag			(a) <i>Ca</i>	(b) <i>Sr</i>
52.	Bone ash contains		[KCET 1992]		(c) <i>Ba</i>	(d) <i>Ra</i>
	(a) <i>CaO</i>	(b) $CaSO_4$		63.	Point out the incor	rect statement regarding Be
	(c) $Ca_3(PO_4)_2$	(d) $Ca(H_2P)$	$(O_4)_2$		(Group-IIA)	
53.	A substance absorbs	$CO_2$ and vi	olently reacts			[MP PMT 1997]
	with water. That subst	ance is			(a) It forms an ionic [AFMC 1988]	c carbide composes on heating
	(a) CaCO <sub>3</sub>	(b) <i>CaO</i>				
	(c) $H_2SO_4$	(d) <i>ZnO</i>			(c) Its halides are co	
<b>-</b> 4	Setting of cement is an		[DPMT 1984]	٠.	(d) It is easily attac	
54.	(a) Exothermic reaction		[DPM1 1964]	64.	-	om rest of the members of its in many ways. The reason for
	(b) Endothermic react				this is its	in many ways. The reason for
	(c) Neither exothermic		mic			[MP PMT 1997]
	(d) None of these				(a) Small size and h	igher electronegativity
55.	Which is quick lime	ı	[EAMCET 1993]		(b) Small size and lo	ower electronegativity
	(a) $Ca(OH)_2$	(b) <i>CaO</i>	5501		(c) Large size and lo	ower ionisation energy
	(c) CaCO <sub>3</sub>	(d) Ca(OH) <sub>2</sub>	+ H . O		(d) Large size and la	argest ionic radius
	-	·	_	65.	The oxide, which is	best soluble in $H_2O$ is [BHU 2001]
56.	A major constituent lime is	of portland c	ement except		(a) $Ba(OH)_2$	(b) $Mg(OH)_2$
	IIIIC 13		[CPMT 1982]		(c) $Sr(OH)_2$	(d) $Ca(OH)_2$
	(a) Silica	(b) Alumina		66.		ne alkaline earth metals that
	(c) Iron oxide	(d) Magnes		00.		atomic number is [BHU 2001]
57.	Portland cement is ma	nufactured by	using [CPMT 198	86]	(a) Ionisation energ	
	(a) Lime stone, clay ar	nd sand	_		(b) Electronegativit	•
	(b) Lime stone, gypsur	n and sand			(c) Solubility of the	ir sulphates
	(c) Lime stone, gypsur	n and alumina	Į.		(d) Solubility of the	ir hydroxides
	(d) Lime stone, clay ar	nd gypsum		67.	In the Alkaline eart	h metals, the element forming
58.	Identify the correct sta	atement [C	BSE PMT 1995]		predominantly cova	lent compound is [BHU 2001]
	(a) Gypsum contains	-	-		(a) <i>Be</i>	(b) <i>Mg</i>
	plaster of calcium	<del>-</del>	=		(c) <i>Sr</i>	(d) <i>Ca</i>
	(b) Gypsum is obtained		-	68.	A mixture of lime pa	aste is sand, water and[RPMT 1997]
- C	(c) Plaster of paris ca	n be obtained	by hydration		(a) Gypsum	(b) Slacked lime
or gy	psum (d) Placton of paris	ia obtaina	l by portial		(c) Quick lime	(d) Lime stone
	(d) Plaster of paris oxidation of gypsu		i by partiai	69.	The formula for cald	cium chlorite is[CBSE PMT 1994, 96]
59.	Which of the follow		es on going		(a) $Ca(ClO_4)_2$	(b) $Ca(ClO_3)_2$
٠,٠	gradually from Be to				(c) CaClO <sub>2</sub>	(d) $Ca(ClO_2)_2$
	(a) Basic character of	=		<b>50</b>	<del>-</del>	ostances gives same gaseous
	(b) Solubility of sulpha	ates in water		70.	_	e react with water[CBSE PMT 1994]

(a) Ca and  $CaH_2$  (b) Na and  $Na_2O_2$ 

(c) K and  $KO_2$  (d) Ba and  $BaO_2$ 

(c) Solubility of hydroxides in water

**60.** Alkaline earth metals are

(d) Strength of elements as reducing agent

[MP PMT 1996]

71.	Magnesium does not decompose the [AFMC 1999]		(c) $BeCl_2 < BaCl_2 < MgCl_2 < CaCl_2$
	(a) Steam (b) Hot water		(d) $BaCl_2 < CaCl_2 < MgCl_2 < BeCl_2$
	(c) Cold Water (d) Semi hot water	83.	$MgCl_2.6H_2O$ when heated gives [CPMT 1997]
<b>72.</b>	Alkaline earth metals are denser than alkali	٥).	
	metals because metallic bonding is		(a) Magnasium diebloride
	(a) Stronger (b) Weaker		<ul><li>(b) Magnesium dichloride</li><li>(c) Magnesium oxide</li></ul>
	(c) Not present (d) Volatile		(d) Magnesium chloride
73•	Property of the alkaline earth metals that increases with their atomic number is [IIT 1997]	84.	Which of the following hydroxide is insoluble in
	(a) Ionisation energy	04.	water
	(b) Solubility of their hydroxides		[AIIMS 2001]
	(c) Solubility of their sulphates		(a) $Be(OH)_2$ (b) $Mg(OH)_2$
	(d) Electronegativity		(c) $Ca(OH)_2$ (d) $Ba(OH)_2$
74.	A metal is burnt in air and the ash on moistening	Q	
	smells of $NH_3$ . The metal is	°5 <sub>[]</sub>	Which of the following statements is false [BHU 2005] (a) $CaOCl_2$ gives $OH^-$ , $Cl^-$ and $OCl^-$ in aqueous
	(a) <i>Na</i> (b) <i>Fe</i>	solu	
	(c) $Mg$ (d) $Al$		(b) Diamond and graphite are allotrops of carbon
75.	Alkaline earth metals come under [Bihar MEE 1996]		(c) Bleaching action of $Cl_2$ in moist condition is
	(a) Halogens (b) Representative		not permanent
elem	nents		(d) Calomel is $Hg_2Cl_2$
elem	(c) Transition elements (d) Inner transition nents	86.	A metal $M$ readily forms its sulphate $MSO_4$ which
	(e) None of these		is water-soluble. It forms its oxide MO which
76.	Which of the following alkaline-earth metal		becomes inert on heating. It forms its insoluble
	hydroxides is the strongest base [CPMT 1996]		hydroxide $M(OH)_2$ which is soluble in $NaOH$
	(a) $Be(OH)_2$ (b) $Mg(OH)_2$		solution. Then $M$ is [AIEEE 2002]
	(c) $Ca(OH)_2$ (d) $Ba(OH)_2$		(a) <i>Mg</i> (b) <i>Ba</i>
77•	Which one of the following is the strongest base	_	(c) Ca (d) Be
	[Pb. PMT 1998]	87.	In the lime (kiln), the reaction
	(a) $Be(OH)_2$ (b) $Mg(OH)_2$		$CaCO_3(s) \rightarrow CO_2(g)$ goes to completion because
	(c) $Al(OH)_3$ (d) $Si(OH)_4$		[Kerala (Engg.) 2002]
78.	Lime stone is [RPMT 1997]		(a) Of high temperature
	(a) $CaO$ (b) $Ca(OH)_2$		(b) $CaO$ is more stable than $CaCO_3$
	(c) Both (a) and (b) (d) None of these		(c) $CO_2$ escapes simultaneously
79.	Which of the alkaline earth metals is strongest		(d) CaO is not dissociated
	reducing agent [MP PMT 1995]	88.	The ionic compound $BaSO_4$ is insoluble in water
	(a) Ca (b) Sr		due to
	(c) <i>Ba</i> (d) <i>Mg</i>		[CPMT 1999]
80.	Plaster of paris hardens by [CPMT 1994]		(a) High lattice energy (b) Low lattice energy
	(a) Giving off $CO_2$ (b) Changing into $CaCO_3$		(c) Low hydration energy (d) Both (a) and (c)
	(c) Uniting with water (d) Giving out water	89.	which is used to reduced the acidity of soil[DPMT 200
81.	Which is not soluble in water [CPMT 1994]		(a) Calcium hydroxide (b) Ammonium sulphate
	(a) $CaCO_3$ (b) $BaCO_3$		(c) Ammonium nitrate (d) Ammonium chloride
	(c) $SrCO_3$ (d) All of these	90.	9 - 1
82.	The correct order of the increasing ionic character		(a) $s$ – block in periodic table
	is		(b) $p$ – block in periodic table
	[MNR 1991; AFMC 1998]		(c) <i>d</i> – block in periodic table
	(a) $BeCl_2 < MgCl_2 < CaCl_2 < BaCl_2$		(d) $f$ – block in periodic table
	(b) $BeCl_2 < MgCl_2 < BaCl_2 < CaCl_2$	91.	The element having atomic number 56 belongs to [AFMC 2002]

#### s and p-Block Elements 767 (a) Actinides (b) Alkaline earth metals (a) $NH_2CONH_2$ (b) $NH_4NO_3$ (c) Transition series (d) Lanthanides (c) CaNCN (d) $KNO_2$ The thermal stability of alkaline earth metal 103. Plaster of paris is used [Pb. CET 2000; CPMT 2000] carbonates $MgCO_3$ , $CaCO_3$ , $BaCO_3$ and (a) In surgery and dentistry SrCO<sub>3</sub> decreases as (b) As a white wash [MP PMT 2002] (c) As a constituent of tooth paste (a) $CaCO_3 > SrCO_3 > MgCO_3 > BaCO_3$ (d) For the preparation of RCC (b) $BaCO_3 > SrCO_3 > MgCO_3 > CaCO_3$ 104. Iron pipes lying under acidic soil are often attached to blocks of magnesium for protection (c) $BaCO_3 > SrCO_3 > CaCO_3 > MgCO_3$ from rusting. Magnesium offers protection to iron (d) $MgCO_3 > CaCO_3 > SrCO_3 > BaCO_3$ against corrosion because it A certain metal M is used to prepare an antacid, [DPMT 2004; BHU 2004] which is used as a medicine in acidity. This metal (a) Prevents air from reaching the surface of iron accidently catches fire which can not be put out (b) is more readily converted into positive ions by using $CO_2$ based extinguishers. The metal M is (c) Is higher than iron (a) Ca (d) Forms a corrosion-resistance alloy with iron (c) Mq (d) All of these **105.** Among K, Ca, Fe, and Zn, the element which can $Be(OH)_2$ is insoluble in water while $Ba(OH)_2$ is 94. form more than one binary compound with highly soluble due to [AMU 2002] chlorine is (a) Bond order (b) Lattice energy [CBSE PMT 2004] difference (a) K (b) Ca (c) Common ion effect (d) Hard acid (c) Fe (d) Zn Which of the following gives a green colour to 95. **106.** *Li* shows the diagonal relationship with [Pb.CET 2001] flame (b) B (a) Mq [AFMC 2001] (c) Al (d) C (a) Barium (b) Calcium **107.** A sodium salt on treatment with $MgCl_2$ gives (c) Strontium (d) None of these white precipitate only on heating. The anion of **96.** Sparingly soluble salt is [RPMT 1999] the sodium salt is (a) KCl (b) NaCl [IIT JEE Screening 2004] (c) NH Cl (d) $BaSO_{A}$ (b) $CO_3^{2-}$ (a) $HCO_3^-$ 97. Among the alkaline earth metals the element forming predominantly covalent compound is [MP PET 1999] (c) $NO_3^-$ (d) $SO_4^{2-}$ (a) Barium (b) Strontium **108.** $MgCl_2.6H_2O$ . When heated gives [MHCET 2003] (c) Calcium (d) Berylium (a) Magnesium oxide **98.** Peroxide bond is present in [RPET 2003] (b) Magnesium oxychloride (a) *MgO* (b) CaO (c) Magnesium dichloride (c) $Li_2O$ (d) $BaO_2$ (d) Magnesium chloride 99. Least ionic character is found in [CPMT 1993] 109. Mg burns in CO to produce [Pb.PMT 2001] (a) Mg (b) Sr (a) $MgO_2$ (b) $MgCO_3$ (d) Ra (c) MgO + CO(d) MgO + C100. The number of water molecules in gypsum and plaster of paris respectively are 110. Sorel [Pbe#M#t1999] [Pb.CET 2003] (a) 1/2 and 2 (b) 2 and 1/2 (a) Portland cement +MgO(c) 2 and 1 (d) 5 and 2 (b) $MgCl_2.CaSiO_3.2H_2O$ 101. Which of the following is formed when calcium (c) CaSiO<sub>3</sub>.MgCO<sub>3</sub> combines with oxygen [MH CET 2000] (a) Ca (b) CaO (d) $MgCl_2.5MgO.xH_2O$ (c) *CaO*<sub>2</sub> (d) $Ca_2O_2$ 111. Colemnite is [AFMC 2004] 102. Slow acting nitrogenous fertilizer among the (a) $Ca[B_3O_4(OH)_2].2H_2O$

[DCE 2003]

(b)  $Ca_2B_6O_{11}.5H_2O$ 

(c)  $Ca(OH)_2$ 

following is

(d)  $Na_2B_4O_7.2H_2O$ 

### **Boron family**

Which of the following statements about  $H_2BO_2$ 1. is not correct [CBSE PMT 1994]

- (a) It is a strong tribasic acid
- (b) It is prepared by acidifying an aqueous solution of borax
- (c) It has a layer structure in which planar  $BO_2$ units are joined by hydrogen bonds
- (d) It does not act as proton donor but acts as a Lewis acid by accepting hydroxyl ion
- The type of hybridisation of boron in diborane is 2. [CPMT 1999]
  - (a) sp-hybridisation
  - (b)  $sp^2$  hybridisation
  - (c)  $sp^3$  hybridisation
  - (d)  $sp^3d^2$  hybridisation
- In the reaction  $B_2O_3 + C + Cl_2 \rightarrow A + CO$ . The A is 3.

[Pb. PMT 2000]

- (a)  $BCl_3$
- (b) *BCl*,
- (c)  $B_2Cl_2$
- (d)  $CCl_2$
- The molecular formula of felspar is [MP PMT 2003] 4.
  - (a)  $K_2O.Al_2O_3.6SiO_2$
- (b)  $K_2O.3Al_2O_3.6SiO_2$
- (c)  $Na_3AlF_6$
- (d)  $CaSO_4.2H_2O$
- The most acidic of the following compounds is 5.

[Bihar CEE 1995]

- (a)  $P_2O_3$
- (b)  $Sb_2O_3$
- (c)  $B_2O_3$
- (d)  $As_2O_3$
- Identify the statement that is not correct as far as 6. structure of diborane is concerned [Pb. PMT 1998]
- (a) There are two bridging hydrogen atoms in diborane
  - (b) Each boron atom forms four bonds in diborane
  - (c) The hydrogen atoms are not in the same plane in diborane
  - (d) All B H bonds in diborane are similar
- Soft heavy metal melts at  $30^{\,o}\,C$  and is used in 7.
  - (a) Galium
- (b) Sodium
- (c) Potassium
- (d) Caesium
- 8. Which of the following is formed when aluminium oxide and carbon is strongly heated in dry chlorine gas

[AFMC 2000]

- (a) Aluminium chloride
- (b) Hydrate aluminium chloride
- (c) Anhydrous aluminium chloride

- (d) None of these
- Which metal burn in air at high temperature with 9. the evolution of much heat
  - (a) *Cu*
- (b) Hg
- (c) Pb
- (d) Al
- Aluminium hydroxide is soluble in excess of sodium hydroxide forming the ion [AMU 2001]
  - (a)  $AlO_2^{+3}$
- (b)  $AlO_{2}^{-3}$
- (c)  $AlO_2^-$
- (d)  $AlO_3^-$
- 11. Boron form covalent compound due to [Pb. PMT 2000]
  - (a) Higher ionization energy
  - (b) Lower ionization energy
  - (c) Small size
  - (d) Both (a) and (c)
- In diborane, the two H-B-H angles are nearly 12.

[AIIMS 2005]

- (a) 60°, 120°
- (b) 95°, 120°
- (c) 95°, 150°
- (d) 120°, 180°
- 13. Which of the following is a non-metal[MP PMT 1999]
  - (a) Gallium
- (b) Indium
- (c) Boron
- (d) Aluminium
- Which of the following is most acidic [BHU 1998]
  - (a)  $Na_2O$
- (b) *MgO*
- (c)  $Al_2O_3$
- (d) CaO
- When orthoboric acid  $(H_3BO_3)$  is heated, the 15. residue left is

[Pb. PMT 2002]

- (a) Metaboric acid
- (b) Boron
- (c) Boric anhydride
- (d) Borax
- 16. Which of the following form dimeric halides

[Roorkee Qualifying 1998]

- (a) *Al*
- (b) Mg

(c) In

- (d) Ga
- The liquid field metal expanding on solidification

[AIIMS 2004]

- (a) Ga
- (b) Al
- (c) Zn
- (d) Cu
- Aluminium chloride exists as dimer,  $Al_2Cl_6$  in solid state as well as in solution of non-polar making heat sensitive thermometers the metal is [RPET 2000] olvents such as benzene. When dissolved in water, it gives [AIEEE 2004]
  - (a)  $[Al(OH)_6]^{3-} + 3HCl$
- (b)  $[Al(H_2O)_6]^{3+} + 3Cl^{-}$
- (c)  $Al^{3+} + 3Cl^{-}$
- (d)  $Al_2O_3 + 6HCl$
- The hardest substance amongst the following is 19. [Kerala PMT 2004]
  - (a)  $Be_2C$
- (b) Graphite
- (c) Titanium
- (d) SiC
- (e)  $B_AC$

(c) Iron undergoes reaction easily with water

Aluminium vessels should not be washed with

materials containing washing soda since[KCET 1993]

(d) Iron forms mono and divalent ions

(a) Washing soda is expensive

over hot aluminium

(b) By passing dry HCl

henzene powder [Pb. CET 2001] (c) By passing dry  $Cl_2$ over hot aluminium (a) Borazine (b) Boron nitride powder (c) p-dichlorobenzene (d) Phosphonitrilic acid (d) By passing dry Cl2 over a hot mixture of Which of the following is only acidic in nature 21. alumina and coke [AIIMS 2004] An element A dissolves both in acid and alkali. It 30. (a)  $Be(OH)_2$ (b)  $Mg(OH)_2$ is an example of [NCERT 1972] (c)  $B(OH)_3$ (d)  $Al(OH)_3$ (b)Dimorphic nature of A (a) Allotropic nature of A 22. Moissan boron is (c) Amorphous nature of A (d)Amphoteric nature of A[DCE 2003] (a) Amorphous boron of ultra purity Hydrogen gas will not reduce [IIT 1984] (b) Crystalline boron of ultra purity (a) Heated cupric oxide (c) Amorphous boron of low purity (b) Heated ferric oxide (d) Crystalline boron of low purity (c) Heated stannic oxide Which of the following does not exist in free form (d) Heated aluminium oxide 23. [Kerala PMT 2004] 32. Conc.  $HNO_3$ (a)  $BF_3$ (b)  $BCl_3$ (a) Reacts with aluminium vigrously (c)  $BBr_3$ (d)  $BH_3$ (b) Reacts with aluminium to form aluminium nitrate (e) None of these (c) Does not react with aluminium 24. Alumina is [DCE 2002] (d) Reacts with platinum (a) Acidic (b) Basic Anhydrous AlCl<sub>3</sub> is obtained from (c) Amphoteric (d) None of these [BHU 1980; CPMT 1982] The most abundant metal in the earth crust is 25. (a) HCl and aluminium metal [Pb. CET 2004] (b) Aluminium and chlorine gas (a) Al (b) Ca (c) Hydrogen chloride gas and aluminium metal (c) Fe (d) Na (d) None of the above 26. Crystalline metal can be transformed into metallic glass by Which is true for an element R present in III 34. [NCERT 1984] group of the periodic table [EAMCET 1991] (a) Alloying (a) It is gas at room temperature (b) It has oxidation state of +4 (b) Pressing into thin plates (c) Slow cooling of molten metal (c) It forms  $R_2O_3$ (d) Very rapid cooling of a spray of the molten (d) It forms  $RX_2$ metal When Al is added to KOH solution 35. Which metal is protected by a layer of its own 27. [NCERT 1974, 76; CPMT 1977] oxide (a) No action takes place [NCERT 1981; DPMT 1983; BHU 1998] (b) Oxygen is evolved (a) *Al* (b) Ag (c) Water is produced (d) Fe (c) Au (d) Hydrogen is evolved **28.** Aluminium is a self-preserving metal, because Aluminium is more reactive than iron. But 36. (a) It is not tarnished by air aluminium is less easily corroded than iron (b) A thin film of basic carbonate on its surface because [KCET 1993] (c) A non-porous layer of oxide is formed on its (a) Aluminium is a noble metal surface (b) Oxygen forms a protective oxide layer (d) It is not affected by salt water

20.

29.

Which of the following is known as inorganic

Anhydrous AlCl<sub>3</sub> cannot be obtained from which

[CPMT 1987]

37.

of the following reactions

(a) Heating  $AlCl_3.6H_2O$ 

- (b) Washing soda is easily decomposed
- (c) Washing soda reacts with aluminium to form soluble aluminate
- (d) Washing soda reacts with aluminium to form insoluble aluminium oxide
- **38.** Which of the statements about anhydrous aluminium chloride is correct [IIT 1981]
  - (a) It exists as AlCl<sub>3</sub> molecule
  - (b) It is not easily hydrolysed
  - (c) It sublimes at  $100 \, ^{o} \, C$  under vacuum
  - (d) It is a strong Lewis base
- 39. Common alum is

#### [DPMT 1982; CPMT 1978; AMU 1982, 83]

- (a)  $K_2SO_4.Al_2(SO_4)_3.24H_2O$
- (b)  $K_2SO_4.Cr_2(SO_4)_3.24H_2O$
- (c)  $K_2SO_4$ .  $Fe_2(SO_4)_3$ .  $24H_2O$
- (d)  $(NH_4)_2 SO_4 .FeSO_4 .6H_2O$
- **40.** Which of the following is not true about potash alum

#### [MNR 1993; UPSEAT 2002]

- (a) Its empirical formula is  $KAl(SO_4)_2.12H_2O$
- (b) Its aqueous solution is basic
- (c) It is used in dyeing industries
- (d) On heating it melts in its water of crystallization
- **41.** Which one of the following is correct statement
  - (a) The hydroxide of aluminium is more acidic than that of boron
  - (b) The hydroxide of boron is basic, while that of aluminium is amphoteric
  - (c) The hydroxide of boron is acidic, while that of aluminium is amphoteric
  - (d) The hydroxide of boron and aluminium are amphoteric
- **42.** AlCl<sub>3</sub> is [AFMC 1995]
  - (a) Anhydrous and covalent (b)Anhydrous and ionic
- acidic

  Aluminium (III) chloride forms a dimer because

(c) Covalent and basic (d) Coordinate

- **43.** Aluminium (III) chloride forms a dimer because

  [CBSE PMT 1995]
  - (a) Higher coordination number can be achieved by aluminium
  - (b) Aluminium has high ionization energy
  - (c) Aluminium belongs to III group
  - (d) It cannot form a trimer
- **44.** Aluminium has a great affinity for oxygen and its oxidation is an exothermic process. This fact is made use of in

#### [MP PMT 1997]

- (a) Preparing thin foils of aluminium
- (b) Making utensils
- (c) Preparing duralumin alloy

- (d) Thermite welding
- 45. Number of water molecules in Mohr's salt is [CPMT 1997; AIIMS 2001; JIPMER 2001]
  - (a) 7

(b) 6

(c) 5

- (d) 8
- **46.** Which of the following is an amphoteric oxide

[BHU 2001]

- (a) MgO
- (b)  $Al_2O_3$
- (c)  $Cl_2O_7$
- (d)  $Ti_2O_2$
- **47.** Aluminium oxide is not reduced by chemical reactions since

#### [KCET 2002]

- (a) Aluminium oxide is reactive
- (b) Reducing agents contaminate
- (c) Aluminium oxide is highly stable
- (d) The process pollutes the environment
- **48.** Aluminium is not used

[DPMT 2002]

- (a) In silvery paints
- (b) For making utensils
- (c) As a reducing agent
- (d) As oxidizer in metallurgy
- **49.** In the thermite process the reducing agent is

[Pb. PMT 2002]

- (a) *Al*
- (b) C
- (c) Mq
- (d) Na
- **50.** In Goldschmidt aluminothermic process, thermite contains

### [KCET 2003]

- (a) 3 parts of  $Al_2O_3$  and 4 parts of Al
- (b) 3 parts of  $Fe_2O_3$  and 2 parts of Al
- (c) 3 parts of  $Fe_2O_3$  and 1 part of Al
- (d) 1 part of  $Fe_2O_3$  and 1 part of Al
- **51.** Bauxite containing impurities of iron oxide is purified by

#### [CPMT 1987; AIIMS 1998]

- (a) Hoop's process
- (b) Serpeck's process
- (c) Baeyer's process
- (d) Electrolytic process
- **52.** In the purification of bauxite by Hall's process
  - (a) Bauxite ore is heated with  $\it NaOH$  solution at  $50^{\it o}\,\it C$
  - (b) Bauxite ore is fused with  $Na_2CO_3$
  - (c) Bauxite ore is fused with coke and heated at  $1800\,^{o}\,C$  in a current of nitrogen
  - (d) Bauxite ore is heated with NaHCO<sub>3</sub>
- **53.** Which one is used as a bye-product in Serpeck's process
  - (a)  $NH_3$
- (b) *CO*<sub>2</sub>
- (c)  $N_2$
- (d)  $PH_3$
- 54. In the metallurgy of aluminium, cryolite is mixed in the molten state because it [Roorkee 1995]
  - (a) Increases the melting point of alumina
  - (b) Oxidises alumina

- (c) Reduces alumina
- (d) Decreases the melting point of alumina
- **55.** In the electrolytic extraction of aluminium, cryolite is used

# [NCERT 1981; CPMT 1989; RPMT 2000; MP PMT 2000, 02]

- (a) To obtain more aluminium
- (b) To decrease temperature to dissolve bauxite
- (c) To protect the anode
- (d) As reducing agent
- **56.** In the extraction of aluminium, bauxite is dissolved in cryolite because
  - (a) It acts as a solvent
  - (b) It reduces melting point of aluminium oxide
  - (c) It increases the resistance of aluminium oxide
  - (d) Bauxite becomes active
- 57. In the extraction of aluminium the electrolyte is [CBSE PMT 1989; AIEEE 2002]
  - (a) Fused cryolite with felspar
  - (b) Fused cryolite with fluorspar
  - (c) Pure alumina in molten cryolite
  - (d) Pure alumina with bauxite and molten cryolite
- 58. Aluminium is obtained by [KCET 1992; RPMT 2002]
  - (a) Reducing  $Al_2O_3$  with coke
    - (b) Electrolysing  $Al_2O_3$  dissolved in  $Na_3AlF_6$
    - (c) Reducing  $Al_2O_3$  with chromium
    - (d) Heating alumina and cryolite
- **59.** In the electrolysis of alumina, cryolite is added to [IIT 1986; BHU 1987]
  - (a) Increase the melting point of alumina
  - (b) Increase the electrical conductivity
  - (c) Minimise the anodic effect
  - (d) Remove impurities from alumina
- **60.** The function of fluorspar in the electrolytic reduction of alumina dissolved in fused cryolite  $(Na_3AlF_6)$  is

[KCET 1993; IIT 1993]

- (a) As a catalyst
- (b) To lower the temperature of the melt and to make the fused mixture very conducting
- (c) To decrease the rate of oxidation of carbon at the anode
- (d) None of the above
- **61.** For purification of alumina, the modern processes most useful when (i) the impurity present is a lot of iron oxides and (ii) the impurity present is a lot of silica, are
  - (a) For (i) Hall's process; for (ii) Baeyer's process
- (b) For (i) Hall's process; for (ii) Serpeck's process
- (c) For (i) Serpeck's process; for (ii) Baeyer's process
  - (d) For (i) Baeyer's process; for (ii) Serpeck's process
- **62.** For the electrolytic production of aluminium, (i) the cathode and (ii) the anode are made of

- (a) (i) Platinum and (ii) Iron
- (b) (i) Copper and (ii) Iron
- (c) (i) Copper and (ii) Carbon
- (d) (i) Carbon and (ii) Carbon
- **63.** In the commercial electrochemical process for aluminium extraction, the electrolyte used is[IIT-JEE 1999]
  - (a)  $Al(OH)_3$  in NaOH solution
  - (b) An aqueous solution of  $Al_2(SO_4)_3$
  - (c) A molten mixture of  $Al_2O_3$  and  $Na_3AlF_6$
  - (d) A molten mixture of AlO(OH) and  $Al(OH)_3$
- **64.** In electrolysis of aluminium oxide which of the following is added to accelerate the process[AFMC 1999; C
  - (a) Silica

(b) Cryolite

(c) Nickel

- (d) Silicate
- 55. The purification of alumina is called

#### [CPMT 1997; AFMC 1998; AIIMS 1999]

- (a) Bosch process
- (b) Caster process
- (c) Baeyer's process
- (d) Hoop's process
- **66.** Electrolytic reduction of alumina to aluminium by Hall-Heroult process is carried out in the presence of

[IIT-JEE (Screening) 2000]

- (a) NaCl
- (b) Fluorite
- (c) Cryolite which forms a melt with lower melting temperature
- (d) Cryolite which forms a melt with higher melting temperature
- **67.** In the electrolytic method of obtaining aluminium from purified bauxite, cryolite is added to the charge in order to

[KCET 2004]

- (a) Minimize the heat loss due to radiation
- (b) Protect aluminium produced from oxygen
- (c) Dissolve bauxite and render it conductor of electricity
- (d) Lower the melting point of bauxite
- **68.** Hoop's process is used for the purification of the metal

[MP PET 1995; MP PMT 2001]

- (a) *Al*
- (b) Zn
- (c) Ag
- (d) Cu
- **69.** Purification of aluminium done by electrolytic refining is known as

[CPMT 1989; CBSE PMT 1999; RPET 2003; BCECE 2005]

- (a) Serpeck's process
- (b) Hall's process
- (c) Baeyer's process
- (d) Hoop's process
- 70. In the Hoope's process for refining of aluminium, the fused materials form three different layers and they remain separated during electrolysis also. This is because

[MP PET 1996]

- (a) The upper layer is kept attracted by the cathode and the lower layer is kept attracted by the anode
- (b) There is special arrangement in the cell to keep the layers separate
- (c) The 3 layers have different densities
- (d) The  $\, 3 \,$  layers are maintained at different temperatures
- **71.** During metallurgy of aluminium bauxite is dissolved in cryolite because
  - (a) Bauxite is non-electrolyte
  - (b) Cryolite is a flux
  - (c) Cryolite acts as an electrolyte
  - (d) All are correct
- **72.** For the electrolytic refining of aluminium, the three fused layers consist of

	<b>Bottom Layer</b>	Middle Layer	Upper Layer
(a)	Cathode of	Cryolite and	Anode of Al
	pure <i>Al</i>	fluorspar	and <i>Cu</i> alloy
(b)	Cathode of <i>Al</i>	Bauxite and	Anode of
	and <i>Cu</i> alloy	cryolite	pure <i>Al</i>
(c)	Anode of Al	Cryolite and	Cathode of
	and <i>Cu</i> alloy	barium	pure <i>Al</i>
		fluoride	
(d)	Anode of	Bauxite,	Cathode of
	impure <i>Al</i>	cryolite and	pure <i>Al</i>
		fluorspar	

- **73.** Heating an aqueous solution of aluminium chloride to dryness will give [AIEEE 2005]
  - (a)  $AlCl_3$
- (b)  $Al_2Cl_6$
- (c)  $Al_2O_3$
- (d)  $Al(OH)Cl_2$
- **74.** The structure of diborane  $(B_2H_6)$  contains[AIEEE 2005]
  - (a) Four 2c-2e bonds and two 3c-2e bonds
  - (b) Two 2c-2e bonds and four 3c-2e bonds
  - (c) Two 2c-2e bonds and two 3c-3e bonds
  - (d) Four 2c-2e bonds and four 3c-2e bonds
- **75.** Which of the following is the electron deficient molecule

[CBSE PMT 2005]

- (a)  $B_2H_6$
- (b)  $C_2H_6$
- (c) PH<sub>3</sub>
- (d) SiH 4
- **76.** In Hall's process, the main reagent is mixed with [AFMC 2005]
  - (a) NaF
- (b)  $Na_3AlF_6$
- (c)  $AlF_3$
- (d) None of these
- 77. Acedic strength of Boron trihalide are in order of [Kerala CET 2005]
  - (a)  $BF_3 < BCl_3 < BBr_3 < BI_3$
  - (b)  $BI_3 < BBr_3 < BCl_3 < BF_3$
  - (c)  $BBr_3 < BCl_3 < BF_3 < BI_3$
  - (d)  $BF_3 < BI_3 < BCl_3 < BBr_3$

### Carbon family

1. Carbon and silicon belong to (IV) group. The maximum coordination number of carbon in commonly occurring compounds is 4, whereas that of silicon is 6. This is due to

[CBSE PMT 1994]

- (a) Large size of silicon
- (b) More electropositive nature of silicon
- (c) Availability of low lying *d*-orbitals in silicon
- (d) Both (a) and (b)
- 2. The ionic carbide is

[JIPMER 2000]

- (a) *ZnC*
- (b) TiC
- (c) SiC
- (d)  $CaC_2$

3.  $PbO_2$  is

[JIPMER 2000]

- (a) Basic
- (b) Acidic
- (c) Neutral
- (d) Amphoteric
- Lead pipes are not suitable for drinking water because

[JIPMER 2000]

- (a) A layer of lead dioxide is deposited over pipes
- (b) Lead reacts with air to form litharge
- (c) Lead reacts with water containing air to form Pb(OH)<sub>2</sub>
- (d) Lead forms basic lead carbonate
- 5. Silicon dioxide is formed by the reaction of

[KCET (Med.) 2001]

- (a)  $SiCl_4 + 2H_2O$
- (b)  $SiO_2 + 4HF$
- (c)  $SiO_2 + NaOH$
- (d)  $SiCl_4 + NaOH$
- **6.** Which alkali metal carbonate decomposes on heating to liberate  $CO_2$  gas
  - (a)  $Li_2CO_3$
- (b)  $CaCO_3$
- (c)  $Na_2CO_3$
- (d)  $Al_2CO_3$
- **7.** Which of the following gives propyne or hydrolysis

[AIIMS 2005]

- (a)  $Al_4C_3$
- (b)  $Mg_2C_3$
- (c)  $B_4C$
- (d)  $La_4C_3$
- **8.** Which one of the following statements is not correct

[CBSE PMT 1994]

- (a) Zinc dissolves in sodium hydroxide solution
- (b) Carbon monoxide reduces iron (III) oxide to

iron

- (c) Mercury (II) iodide dissolves in excess of potassium iodide solution
- (d) Tin (IV) chloride is made by dissolving tin solution in concentrated hydrochloric acid

**9.** In laboratory silicon can be prepared by the reaction

[Pb. PMT 1999]

- (a) By heating carbon in electric furnace
- (b) By heating potassium with potassium dichromate
- (c) Silica with magnesium
- (d) None of these
- 10. Which of the following is the correct statement for red lead

[AIIMS 2000]

- (a) It is an active form of lead
- (b) Its molecular formula is  $Pb_2O_3$
- (c) It decomposes into Pb and  $CO_2$
- (d) It decomposes into PbO and  $O_2$
- 11. Suppose you have to determine the percentage of carbon dioxide in a sample of a gas available in a container. Which is the best absorbing material for the carbon dioxide

[Pb. PMT 2001]

- (a) Heated copper oxide
- (b) Cold, solid calcium chloride
- (c) Cold, solid calcium hydroxide
- (d) Heated charcoal
- 12. The number and type of bonds between 2 carbon atoms in  $CaC_2$  [UPSEAT 2001]
  - (a) One sigma  $(\sigma)$  and one  $pi(\pi)$  bond
  - (b) One sigma  $(\sigma)$  and two pi  $(\pi)$  bond
  - (c) One sigma  $(\sigma)$  and half pi  $(\pi)$  bond
  - (d) One sigma  $(\sigma)$  bond
- 13. Metalloid among the following is [DPMT 2001]
  - (a) *Si*

(b) C

[Pb-PMH,2000]

(d) *Ge* 

14. 'Lead pencil' contains

- [DPMT 2001; IIT 1990]
- (a) *PbS*
- (b) Graphite
- (c) FeS
- (d) *Pb*
- **15.** Nitrogen gas is absorbed by
  - by [DPMT 2001]
  - (a) Calcium hydroxide
- (b) Ferrous sulphate
- (c) Calcium carbide
- (d) Aluminium carbide
- **16.** In laboratory silicon can be prepared by the reaction

[Pb. PMT 1999; AFMC 2002]

- (a) Silica with magnesium
- (b) By heating carbon in electric furnace
- (c) By heating potassium fluosilicate with potassium
- (d) None of these
- 17. Formation of in-numberable compounds of carbon is due to its
  - (a) High reactivity
  - (b) Catenation tendency

#### 772 s and p-Block Elements (c) Covalent and ionic tendency Which gas is used in excess water 29. [BVP 2003] (a) *CO*<sub>2</sub> (d) Different valency (b) SO<sub>2</sub> Colour is imported to glass by mixing [Pb. PMT 2002] 18. (c) CO (d) Water vapours (a) Synthetic dyes (b) Metal oxide The compound which does not possess a peroxide 30. linkage is (c) Oxides of non-metal (d) Coloured salt (b) CrO<sub>5</sub> (a) $Na_2O_2$ Which of the following is insoluble in water [MP PET 2002] (c) $H_2SO_5$ (d) $PbO_2$ (a) $Na_2CO_3$ (b) CaCO<sub>3</sub> Silicon is an important constituent of [MH CET 2001] (c) $ZnCO_3$ (d) $Al_2(CO_3)_3$ 31. (a) Rocks (b) Amalgams 20. In which of the following the inert pair effect is (c) Chlorophyll (d) Haemoglobin most prominent [MP PMT 2000] 32. Carborundum is (b) Si (a) C [AFMC 2002; MH CET 2003; BHU 2003, 05] (c) *Ge* (d) Pb (a) SiC (b) AlCl<sub>3</sub> Plumbosolvancy implies dissolution of lead in [DCE 1999] (a) Bases (b) Acids (d) $Al_2O_3.2H_2O$ (c) $Al_2(SO_4)_3$ (c) Ordinary water (d) CuSO sol 33. $SiF_4$ gets hydrolysed giving ...... [Orissa JEE 2002] Which of the following glass is used in making (b) Si(OH), F, (a) $SiO_2$ wind screen of automobiles [AIIMS 1999; Pb. CET 2000] (a) Crook's (b) Jena (c) $H_2SiF_6$ (d) $Si(OH)_A$ (c) Safety (d) Pyrex Glass is a [AIEEE 2003; RPET 2003] **23.** Glass reacts with *HF* to produce (a) Micro-crystaline solid [KCET 2000; CBSE PMT PMT 2000] (b) Super cooled liquid (a) $SiF_4$ (b) $H_2SiF_6$ (d) $Na_3AlF_6$ (c) $H_2SiO_3$ (d) Polymeric mixture The type of glass used in making lenses and 24. $H_2O_2$ on reaction with *PbS* gives [RPET 2003] 35. prisms is (a) *PbO* (b) $PbSO_A$ [JIPMER 1999] (d) PbHSO 4 (c) $PbO_2$ (a) A flint glass (b) Jena glass Soldiers of Napolean army while at Alps during (c) Pyrex glass (d) Quartz glass 36. freezing winter suffered a serious problem as When carbon monoxide is passed over solid 25. regards to the tin buttons of their uniforms. caustic soda heated to 200° C, it forms [KCET (Med.) 1999] White metallic tin buttons got converted to grey (a) $Na_2CO_3$ (b) NaHCO<sub>3</sub> powder. This transformation is related to (c) H-COONa(d) CH<sub>3</sub>COONa [AIEEE 2004] 26. Which is used to produce smoke screens [AFMC 2005] (a) A change in the partial pressure of oxygen in the air (a) Calcium phosphide (b) Zinc sulphide (b) A change in the crystalline structure of tin (c) Sodium carbonate (d) Zinc phosphide (c) An interaction with nitrogen of the air at very Sodium oxalate on heating with conc. $H_2SO_4$ 27. low to temperatures gives (d) An interaction with water vapour contained in [Roorkee 2000] the humid air (a) CO only (b) CO, only 37. Solid $CO_2$ is known as dry ice, because[Pb. CET 2000] (c) CO and $CO_2$ (d) $SO_2$ and $SO_3$ (a) It melts at 0°C 28. Extraction of lead by reduction methods is done (b) It evaporates at 40°C (c) It evaporates at $-78^{\circ}C$ without melting [AMU 2000] (d) Its boiling point is more than 199°C (a) Adding more galena into reverberatory furnace Which one of the following statements about the 38. (b) Adding more lead sulphate into reverberatory zeolites is false furnace [CBSE PMT 2004] (c) Adding more galena and coke into the (a) Zeolites are aluminosilicates having three reverberatory furnace dimensional network (d) Self reduction of oxide from sulphide present (b) Some of the $SiO_4^{-4}$ units are replaced by $AlO_4^{-5}$ in the furnace and $AlO_6^{9-}$ ions in zeolites

	(c) They are used as cat		50.	Red lead is	[CPMT 1972, 74, 94; MNR 1985;
	(d) They have open struto take up small mol	icture which enables them			02; Bihar CEE 1995; MP PET 1995]
39.	-	ecules cuts ultraviolet rays[AFMC 20	004]	(a) $Pb_3O_4$	(b) <i>PbO</i>
29.	(a) Soda glass	(b) Crooke's glass	004]	(c) $PbO_2$	(d) $Pb_4O_3$
	(c) Pyrax	(d) None of these	51.	White lead is	[CPMT 1983, 93, 2002; MNR 1984;
40.	•	ium) shows +1 oxidation			MT 1995; UPSEAT 1999; DCE 2000]
10.		nbers show +3 oxidation		(a) $PbCO_3$	(b) PbCO <sub>3</sub> .PbO
	state. Why			(c) $2PbCO_3.Pb(OH)$	$_{2}$ (d) $_{2}PbSO_{4}.PbO$
		[JEE Orissa 2004]	52.	Lead pipes are cor	roded quickly by [AFMC 1981]
	(a) Presence of lone pair	r of electron in <i>Tl</i>		(a) Dil. $H_2SO_4$	(b) Conc. $H_2SO_4$
	(b) Inert pair effect	_		(c) Acetic acid	(d) Water
	(c) Large ionic radius of	f Tl ion	53.	In silicon dioxide	[AIEEE 2005]
	(d) None of these			(a) Each silicon	atom is surrounded by four
41.	Carbon suboxide $C_3O_2$ h	nas [DCE 2003]			s and each oxygen atom is
	(a) Linear structure			bonded to two	
	(b) Bent structure				atom is surrounded by two
	(c) Trigonal planar stru				s and each oxygen atom is so silicon atoms
	(d) Distorted tetrahedra		_		bonded to two oxygen atoms
<b>42.</b>	=	s a mixed oxide[Pb. CET 200	31		ble bonds between silicon and
	(a) $Fe_2O_3$	(b) <i>PbO</i> <sub>2</sub>		oxygen atoms	
	(c) $Pb_3O_4$	(d) <i>BaO</i> <sub>2</sub>	54.	Litharge is chemic	ally [DPMT 1984; JIPMER 2001]
43.	Noble gases are absorbe	d on [BVP 2004]		(a) <i>PbO</i>	(b) <i>PbO</i> <sub>2</sub>
	(a) Anhydrous CaCl <sub>2</sub>	(b) Charcoal		(c) $Pb_3O_4$	(d) $Pb(CH_3COO)_2$
	(c) Conc. $H_2SO_4$	(d) Coconut			-
44.	Lapis lazuli is	[AFMC 2004]	55.		configuration is of group
	(a) Ferrous sulphate	(b) Copper sulphate		(a) IV	(p) III
	(c) Sodium alumino silio	cate (d) Zinc sulphate	56.	(c) V Which of the follo	(d) II wing compounds of elements in
<b>45</b> .	_	statement is correct with	50.		ou expect to be most ionic in
		of elements in the carbon	CET on	character	1
	(a) Atomic size decrease	in atomic number, their[Pb.	CET 20	002]	[NCERT 1978]
	(b) Ionization energy in			(a) <i>CCl</i> <sub>4</sub>	(b) SiCl <sub>4</sub>
	(c) Metallic character de			(c) <i>PbCl</i> <sub>2</sub>	(d) $PbCl_4$
	(d) Stability of +2 oxida		57.	Which of the follo	wing compounds of lead is used
46.	When tin is treated with	concentrated nitric acid[ <b>D</b> (	CE 2004	in match industry	
	(a) It is converted into s			(a) <i>PbO</i>	(b) <i>PbO</i> <sub>2</sub>
	(b) It is converted into s	stannic nitrate		(c) <i>PbCl</i> <sub>2</sub>	(d) None of these
	(c) It is converted into r	netastannic acid	58.	Type metal is an a	lloy of $Pb$ , $Sb$ and $Sn$ . It consists
	(d) It becomes passive			of	
47.	Solder is an alloy of	[Pb. CET 2003]		(a) Equal amounts	of the three metals
	(a) $Pb + Zn + Sn$	(b) $Pb + Zn$		(b) More amount of	of lead
48.	(c) $Pb + Sn$ A metal used in storage	(d) $Sn + Zn$		(c) More amount of	of antimony
40.	(a) Copper	(b) Lead		(d) More amount of	of tin
	(c) Tin	(d) Nickel	59.	Which is correct of	xidation state of lead[AFMC 1987]
49.		of silicates in which three		(a) + 2, + 4	(b) + 1, + 2
	oxygen atoms of $[SiO_4]^4$			(c) $+ 3, + 4$	(d) + 4
	(a) Pyrosilicate		60.	J	
	(b) Sheet silicate			(a) $2PbSO_4.PbO$	(b) $(CH_3COO)_2Pb$
	(c) Linear chain silicate			(c) $PbCO_3$	(d) $PbCO_3.Pb(OH)_2$
	(d) Three dimensional s	ilicate			

[AIIMS 1982, 83, 87; BHU 1998; CBSE PMT 2000;

	· · · · · · · · · · · · · · · · · · ·						
61.		ving compounds has peroxide		Pb. PMT 2000; AFMC 198	8, 92; MP PET	1997, 2000, 01]	
	linkage			(a) $N_2O$	(b) <i>NO</i>		
	(a) DL ()	[CPMT 1988]		(c) $N_2O_3$	(d) $N_2O_5$		
	(a) $Pb_2O_3$	(b) $SiO_2$	4.	Metaphosphoric acid h	as the formul	a[CPMT 1973, 89, 93	
	(c) <i>CO</i> <sub>2</sub>	(d) $PbO_2$	-	(a) $H_3PO_4$	(b) <i>HPO</i> <sub>3</sub>		
62.	Percentage of lead i	=		(c) $H_2PO_3$	(d) $H_3PO_2$		
	(a) Zero	[CBSE PMT 1999] (b) 20	_				
	(a) Zero (c) 80	(d) 70	5.	Which of the following agent for ammonia gas		suitable drying	
63.		ing has most density		•		89; DPMT 1982;	
٠,٠	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	[CPMT 1996]				1986, 96; 2001;	
	(a) Fe	(b) <i>Cu</i>				reening) 2000]	
	(c) B	(d) <i>Pb</i>		(a) Calcium oxide		_	
64.	Red lead in an exam	_		(b) Anhydrous calcium	chloride		
	( ) 7	[JIPMER 2001]		(c) Phosporus pentoxi			
	(a) Basic	(b) Super		(d) Conc. sulphuric aci			
65.	(c) Mixed	(d) Amphoteric	6.	Each of the following		white and red	
05.	Which of the following lead oxides is 'Sindhur' [MP PET 2002]			phosphorus except tha			
	(a) <i>PbO</i>	(b) <i>PbO</i> <sub>2</sub>		(a) Are both soluble in $CS_2$			
	(c) $Pb_2O_3$	(d) $Pb_3O_4$		(b) Can be oxidised by heating in air			
66.		e phenomenon of allotropy is		(c) Consists of same k	ind of atoms		
	Ziement mowing th	[MP PMT 1999]		(d) Can be converted i	nto one anoth	er	
	(a) Aluminium	(b) Tin	7•	Which of the following is a tetrabasic acid[CPMT 1988]			
	(c) Lead	(d) Copper		(a) Orthophosphorus acid (b)Orthophosphoric acid			
67.	Which of the following element is a metalloid			(c) Metaphosphoric acid (d) Pyrophosphoric acid			
-	[CPMT 2004]		8.	Phosphine is prepared	by the reaction	on of	
	(a) Bi	(b) Sn				PET/PMT 1988]	
	(c) Ge	(d) <i>C</i>		(a) $P$ and $H_2SO_4$	(b) <i>P</i> and	NaOH	
68.	Which gas is liberat	ed when $Al_4C_3$ is hydrolysed		(c) $P$ and $H_2S$	(d) $P$ and	$HNO_3$	
		[AFMC 2005]	9.	Which of the following	g is not known	[MP PET/PMT 1988;	
	(a) $CH_4$	(b) $C_2H_2$		NCERT 1982; (	CBSE PMT 1989	; MP PET 1993]	
	(c) $C_2H_6$	(d) $CO_2$		(a) <i>NCl</i> <sub>5</sub>	(b) $NI_3$		
69.	Which of the follow	ing attacks glass		(c) $SbCl_3$	(d) $NCl_3$		
		[NCERT 1976; AFMC 2005]	10.	Chemical formula for t	he phosphoru	s molecule is	
	(a) HCl	(b) HF		[CPMT 19	976, 80, 84, 90	; BHU 1984, 86;	
	(c) HI	(d) <i>HBr</i>		( ) 5	<i>a</i> > 5	NCERT 1977]	
	Nitro	gen family		(a) <i>P</i>	(b) $P_4$		
	Mitro	gen ranning		(c) $P_2$	(d) $P_5$		
1.	Which of the follow	wing elements does not form	11.	White phosphorus $(P_4)$	) has	[IIT 1998]	
1.	stable diatomic mole		[	ckse sixt <sup>P</sup> 1989;ingle bon	ıds		
	(a) Iodine	(b) Phosphorus		(b) Four <i>P</i> - <i>P</i> single bo			
	(c) Nitrogen	(d) Oxygen		(c) Four lone pairs of	electrons		
2.	•	exture of[ <b>DPMT 1982; CPMT 1978</b> ]		(d) <i>PPP</i> angle of $60^{\circ}$			
	(a) $CO$ and $N_2$	(b) $CO_2$ and $H_2$	12.	Ammonium nitrate dec	_	_	
	(c) $CO$ and $H_2$	(d) $CO_2$ and $N_2$		[NCERT 19	974,75; CPMT 1	973, 78, 88, 94;	
_	-			(a) Ammonia and nitri	c acid	AMU 1984]	
3.	ions to form a brown	ollowing combines with Fe(II)		(b) Nitrous oxide and			
	ions to form a brow.	n complex		(b) INITIOUS OXIUE dilu	water		

(c) Nitrogen, hydrogen and ozone

78, 81

	(d) Nitric oxide, nitroge	n dioxide and hydrogen	23.	On heating a mixture of	of NH <sub>4</sub> Cl and KNO <sub>2</sub> we get
13.	In Birkeland-Eyde process, the raw material used				[CPMT 1972, 79; NCERT 1977]
	is	<b>.</b>		(a) $NH_4NO_3$	(b) $N_2$
	(	[CPMT 1982, 86]		(c) $N_2O$	(d) <i>NO</i>
	(a) Air	(b) $NH_3$	24	_	ng oxide of nitrogen is the
	(c) <i>NO</i> <sub>2</sub>	(d) $HNO_3$	24.	anhydride of $HNO_3$	ig oxide of introgen is the
14.	-	trates, Lead nitrate, Silver n nitrate; the one that		[CPMT 1979,80,89,97; N	MP PET/PMT 1988; KCET 1991;
		ving any solid residue is[NCE	RT 198	CBSE PMT 1989, 91,9	9; EAMCET 1991; NCERT 1975;
	(a) Lead nitrate	(b) Ammonium nitrate		( ) 170	MP PET 1989; MP PMT 1994]
	(c) Silver nitrate	(d) Sodium nitrate		(a) NO	(b) $N_2O_3$
15.	Of the different allotrop	oes of phosphorus, the one		(c) $N_3 O_4$	(d) $N_2O_5$
		s [CPMT 1983; NCERT 1978;	25.	-	factured by heating in a
		999; Kurukshetra CEE 1998]			ture of <b>NCERT 1977; CPMT 1974</b>
	(a) Violet phosphorus	(b) Scarlet phosphorus		(a) Bone ash and coke	
	(c) Red phosphorus	(d) White phosphorus		(b) Bone ash and silica	
16.	Phosphine is generally p	repared in the laboratory		(c) Bone ash, silica an	d coke
		[CPMT 1983, 2003]		(d) None of these	
hvdr	(a) By heating phosp ogen	horus in a current of	26.		ns a solid oxide which when rms an acidic solution, the
5	=	phosphorus with aqueous		element is	inis an acidic solution, the
	solution of caustic p				[CPMT 1972, 78]
	(c) By decomposition of	$P_2H_4$ at 110 ° C		(a) Argon	(b) Potassium
		sphorus with an aqueous		(c) Phosphorus	(d) Sulphur
	solution of caustic se		27.	Dissociation of $H_3PO_4$	occurs in following stages
17.	Which of the following e	elements is most metallic			[CPMT 1976]
		[CPMT 1983; MP PMT 1993]		(a) 1	(b) 2
	(a) Phosphorus	(b) Arsenic		(c) 3	(d) 4
	(c) Antimony	(d) Bismuth	28.	Nitrogen forms how m	any oxides
18.	The basicity of orthopho	sphoric acid is[CPMT 1984, 91	]	(a) 3	(b) 4
	(a) 2	(p) 3		(c) 5	(d) 6
	(c) 4	(d) 5	29.	The <i>P-P-P</i> bond angle i	n white phosphorus is
19.	$HNO_2$ acts as	[AFMC 1992]			[MP PET 1991]
	(a) Oxidising agent	(b) Reducing agent		(a) 120°	<b>(b)</b> 109 ° 28′
	(c) Both (a) and (b)	(d) Its solution is stable		(c) 90°	(d) 60°
20.	Nitrogen dioxide cannot	be obtained by heating	30.		
	[СРМТ	1989; IIT 1985; CPMT 1993]			BSE PMT 1993; MP PMT 1993]
	(a) $KNO_3$	(b) $Pb(NO_3)_2$		(a) Chromium oxide at	
	(c) $Cu(NO_3)_2$	(d) $AgNO_3$		<ul><li>(b) Chromic acid and r</li><li>(c) Chromium oxide and r</li></ul>	_
21.		s passed over <i>CuO</i> gas		(d) Chromic acid and a	_
	evolved is	1 2.2- 2 2	31.		nitric acid is heated, it
		[BCECE 2005]		decomposes to give	
	(a) $N_2$	(b) $N_2O$		(a) $O_2$ and $N_2$	(b) <i>NO</i>
	(c) $HNO_3$	(d) $NO_2$		(c) $N_2O_5$	(d) $NO_2$ and $O_2$
22	_	_	32.		sches fire in air at $30^{\circ} C$ and
22.	Non-combustible hydrid				is[BHU 1973; MP PET 1989,99]
	(a) $NH_3$	(b) <i>PH</i> <sub>3</sub>		<ul><li>(a) Calcium</li><li>(c) Phosphorus</li></ul>	<ul><li>(b) Sodium</li><li>(d) Zinc</li></ul>
	(c) $AsH_3$	(d) $SbH_3$	33.	A solution of ammonia	
			55.		

	//O s and p-bio	CK LIEITICITS			
	( ) ***	[CBSE PMT 1991]	gung		nary calcium phosphate and
	(a) H <sup>+</sup>		gyps	cum (c) Normal calcium pl	noenhate
	(b) <i>OH</i>			(d) Soluble calcium ph	_
	(c) Only $NH_4^+$		44.	_	is allowed to react with
	(d) $OH^-$ , $NH_4^+$ and $N$	•	11.	sufficient quantity of	NaOH , the product obtained
34.		of V A group (N, P, As, Sb and		is	[DPMT 1983; MP PMT 1983]
		lowing properties shows an on from nitrogen to bismuth[C	PMT 108	B <b>a</b> Th) NaHPO.	(b) $Na_2HPO_4$
	(a) Stability of +3 oxid		- 1111 - 131	3	(d) $Na_3PO_4$
	(b) Reducing characte			(c) $NaH_2PO_4$	
	(c) Electronegativity		45.	White phosphorus con	ttains E <b>T (Med.) 2000; MP PET 1990]</b>
	(d) Acidic nature of th	=		(a) $P_5$ molecules	(b) $P_4$ molecules
35.	The important method is	d for the fixation of nitrogen		(c) $P_6$ molecules	•
	15	[CPMT 1985, 94]	46.	*	tion of ammonia an oxide is
	(a) Haber	(b) Solvay	40.	-	in the preparation of $HNO_3$ .
	(c) Deacon	(d) Fischer method		This oxide is	1 1
36.	Which of the following				1984; KCET 1990; AIIMS 1996]
	(a) White phosphorus	[AFMC 1987; KCET 1991] (b) CH <sub>4</sub>		(a) $N_2O_5$	(b) $N_2O_4$
		· ·		(c) NO <sub>2</sub>	(d) <i>NO</i>
	(c) $H_2O$	(d) NaCl	47.	Nitric acid oxidises ph	osphorus to
37•		ogen is prepared by heating		-	[CPMT 1984; JIPMER 2002]
	<ul><li>(a) Calcium cyanamid</li><li>(c) Ammonium hydrox</li></ul>			(a) $H_2 P_2 O_7$	(b) $H_3PO_3$
38.	Nitrous oxide	vide (d)Aiiiiioiiidiii iiitiite		(c) $P_2O_5$	(d) $H_3 PO_4$
30.	(a) Is a mixed oxide		48.	Which one of the follo	owing statements is true for
	(b) Is an acidic oxide			$HNO_2$	
	(c) Is highly soluble in	n hot water			[CPMT 1980, 84]
	(d) Supports the comb	<del>-</del>		(a) It is very stable in	_
39.		g represents laughing gas	redu	(b) it cannot act bot	th as an oxidant and as a
		nipal MEE 1995;MP PMT 1990; 1995; RPMT 1999; AFMC 2002]	1044	(c) It cannot act as an	oxidising agent
	(a) <i>NO</i>	(b) $N_2O$		(d) It cannot act as re	
	(c) $NO_2$	(d) $N_2O_3$	49.	Which oxide is alkalin	e [MP PET 1990]
40.	-	e is proved by the first that		(a) $P_2O_3$	(b) $Bi_2O_3$
40.	with $NaOH$ , it forms	e is proved by the mot that		(c) $As_2O_3$	(d) $B_2O_3$
	(a) Nitrites salt		50.	Which acid is formed	by $P_2O_3$ [MP PET 1991]
	(b) Nitrates salt			(a) $H_3PO_4$	(b) $H_3PO_3$
	(c) Mixture of nitrate	and nitrite		(c) HPO <sub>3</sub>	(d) $H_4 P_2 O_7$
	(d) Ammonia		51.	Which nitrogen trihali	· <del>-</del> ·
41.		ing metal produces nitrous	3	-	kshetra CEE 1998; CPMT 1999]
	oxide with dil. HNO <sub>3</sub>			(a) $NF_3$	(b) NCl <sub>3</sub>
	(a) Fe	(b) Zn		(c) $NBr_3$	(d) $NI_3$
40	(c) Cu	(d) Ag	52.	Dehydrated phosphoru	us trichloride in water gives
42.	form	ing acid exist in polymeric			[MP PET 1990]
	(a) $HPO_3$	(b) $H_4 P_2 O_7$		(a) $HPO_3$	(b) $H_3PO_4$
	(c) $H_3PO_4$	(d) None of these		(c) $H_3PO_2$	(d) $H_3PO_3$
43.	Superphosphate of lin		53.		the manufacture of safe
70.		mal calcium phosphate and		matchsticks	[DDMT 1002 CDMT 10= /1
gyps		1 11 11 1111		(a) White phosphorus	[DPMT 1982, CPMT 1974, 75] (b) Sulphur
				(c) Red phosphorus	(d) Selenium

54.	Which oxide of nitrogen			(d) <i>CaC</i> <sub>2</sub>	
		987; Kurukshetra CEE 1998	<sup>]</sup> 66.	On adding water to	o $BiCl_3$ solution in $HCl$ , the
	(a) $N_2O$	(b) <i>NO</i>		compound produced	
	(c) $N_2O_5$	(d) <i>NO</i> <sub>2</sub>		(a) $Bi_2O_3$	(b) $Bi(OH)_3$
55.	Which oxide do not act	as a reducing agent [MP PET 1990]	1	(c) BiOCl	(d) BiOCl <sub>2</sub>
	(a) NO	(b) NO2	67.	V-A group precipitat	te was dissolved in $HNO_3$ and
	(c) $N_2O$	(d) $N_2O_5$		treated with excess	of $NH_4OH$ . It gives a white
56.	=	ation number of nitrogen	1	ppt. because of	
50.	will be	ition number of introger		(a) $Cu(OH)_2$	(b) $Cd(OH)_2$
	WIII DC	[MP PET 1990	]	(c) $Bi(OH)_3$	(d) $Hg(OH)_2$
	(a) + 3	(b) + 5	_	N, P, As, Sb, Bi elem	ents belong to [DPMT 1982]
	(c) $- 3$ and $+ 3$	(d) + 3 and + 5		(a) VA group	(b) IVA group
57•		the oxidation state o	f	(c) VIIA group	(d) VB group
	phosphorus is+4	[MP PET 1991	69.		llowing elements occur free in
	(a) $P_4O_{11}$	(b) $P_4O_8$	J	nature	
	(c) $P_4O_6$	(d) $H_3PO_4$		(a) Nitrogen	[CPMT 1988] (b) Phosphorus
58.		the oxidation state o	f	(c) Arsenic	(d) Antimony
50.	nitrogen is -1	the oxidation state o	70.		ing elements of group VA does
		[MP PMT 1989]	•	not show allotropy	[CPMT 1980]
	(a) <i>NO</i>	(b) $N_2O$		(a) N	(b) <i>Bi</i>
	(c) $NH_2OH$	(d) $N_2H_4$		(c) P	(d) As
59.	Which of the following	oxide is least acidic	71.	Which does not form	-
		P PMT 1990; CBSE PMT 1996	]	(a) <i>N</i>	(b) <i>P</i>
	(a) $P_4O_6$	(b) $P_4 O_{10}$	72	(c) As The strongest base i	(d) Bi
	(c) $As_4O_6$	(d) $As_4O_{10}$	72.	_	s 39; CPMT 1997; MP PET 2001,03]
60.		hydrides of the <i>V</i> -group	•	(a) $NH_3$	(b) <i>PH</i> <sub>3</sub>
		he order [CBSE PMT 1996]	J	(c) $AsH_3$	(d) SbH <sub>3</sub>
	(a) $SbH_3 > PH_3 > AsH_3$		73.	The most stable hyd:	3
	(b) $NH_3 > SbH_3 > PH_3 >$		/3.	(a) $NH_3$	(b) <i>PH</i> <sub>3</sub>
	(c) $NH_3 > PH_3 > AsH_3 >$	-		(c) $AsH_3$	(d) $SbH_3$
	(d) $SbH_3 > AsH_3 > PH_3$		74	-	st boiling point[CBSE PMT 1989]
61.	Which is least stable	[MP PET 1989]	] /	(a) $NH_3$	(b) <i>PH</i> <sub>3</sub>
	(a) $BiH_3$	(b) <i>SbH</i> <sub>3</sub>		(c) $AsH_3$	(d) $SbH_3$
	(c) $AsH_3$	(d) $PH_3$	75.	-	xplosive[BHU 1984; Roorkee 1989
62.	•	is not hydrolysed[DPMT 20	05] /5		AIIMS 1996; MP PMT 1985, 2001]
	(a) $AsCl_3$	(b) <i>PF</i> <sub>3</sub>		(a) $NCl_3$	(b) <i>PCl</i> <sub>3</sub>
	(c) $SbCl_3$	(d) $NF_3$		(c) $AsCl_3$	(d) All of these
63.		e is maximum for[MP PET	<sup>1990]</sup> 76.	Of the following, the	e most acidic is [EAMCET 1980]
	(a) $AsH_3$	(b) $NH_3$		(a) $As_2O_3$	(b) $P_2O_3$
	(c) $PH_3$	(d) $SbH_3$		(c) $Sb_2O_3$	(d) $Bi_2O_3$
64.		is kept in water?[BCECE 20	05] 77.	Of the following, no	n-existent compound is
	(a) White phosphorous				[NCERT 1975, 79]
<b>~</b> -	(c) Potassium	(d) Calcium	_	(a) $PH_4I$	(b) $As_2O_3$
65.	fertilizer	g substances is used as a	a	(c) SbCl <sub>2</sub>	(d) $As_2H_3$
	(a) $K_2SO_4.Al_2(SO_4)_3.24I$	$H_2O$	78.	Pure $N_2$ gas is obtain	ined from [CBSE PMT 1991]
	(b) $Ca(H_3PO_4)_2H_2O + Ca(H_3PO_4)_2H_3O + Ca(H_3PO_4)_2H_3O + Ca(H_3PO_4)_3H_3O + $			(a) $NH_3 + NaNO_2$	(b) $NH_4Cl + NaNO_2$
	(c) $NaAlO_2$	4		(c) $N_2O + Cu$	(d) $(NH_4)_2 Cr_2 O_7$
	(C) 1400 10 2		79.	Pure nitrogen can be	e prepared from

#### 778 s and p-Block Elements [KCET 1991; AFMC 1993; AMU 1985] (b) NO (a) CO (a) $NH_{\perp}OH$ (b) $Ca_3N_2$ (c) ClO<sub>2</sub> (d) $N_2O_5$ On strongly heating $Pb(NO_3)_2$ crystals, the gas (c) $NH_4NO_2$ (d) $Ba(NO_3)_2$ 90. formed is 80. Nitrogen combines with metals to form [CPMT 1981, 93] [NCERT 1980; CPMT 1997] (a) Nitrites (b) Nitrates (a) $NO_2$ (b) $O_2$ (c) Nitrosyl chloride (d) Nitrides Nitrogen is relatively inactive element because (c) $NO_2 + O_2$ (d) NO [CBSE PMT 1992] Nitrogen dioxide is released by heating [AFMC 1992] (a) Its atom has a stable electronic configuration (a) $Pb(NO_3)_2$ (b) $KNO_3$ (b) It has low atomic radius (c) NaNO<sub>2</sub> (d) NaNO 3 (c) Its electronegativity is fairly high Nitric oxide is prepared by the action of $HNO_3$ on (d) Dissociation energy of its molecule is fairly high [AFMC 1990] (a) Fe (b) Cu The cyanide ion, $CN^-$ and $N_2$ are isoelectronic. 82. (d) Sn (c) Zn But in contrast to $CN^-$ , $N_2$ is chemically inert 93. When lightning flash is produced, which gas may because of form [IIT 1992] [EAMCET 1992; AFMC 1989] (a) Low bond energy (a) Nitrous oxide (b) Nitrogen dioxide (b) Absence of bond polarity (c) Dinitrogen pentoxide (d) Nitric oxide (c) Unsymmetrical electron distribution Of the following, which has three electron bond in 94. (d) Presence of more number of electrons in its structure bonding orbitals [CPMT 1986] Which statement is not correct for nitrogen[AIIMS 1991] (a) Nitrous oxide (b) Nitric oxide (a) It has a small size (c) Dinitrogen trioxide (d) Nitrogen pentoxide (b) It does not readily react with $O_2$ Which of the following oxides of nitrogen is 95. (c) It is a typical non-metal neutral (d) *d*-orbitals are available for bonding [CPMT 1988] (a) $N_2O_5$ (b) $N_2O_3$ The element which is essential in nitrogen 84. fixation is (c) $N_2O_4$ (d) $N_2O$ [NCERT 1981] **96.** Oxidation of *NO* in air produces [KCET 1992] (a) Zinc (b) Copper (a) $N_2O$ (b) $N_2O_3$ (c) Molybdenum (d) Boron (d) $N_2O_5$ (c) $NO_2$ **85.** Laughing gas is prepared by heating[EAMCET 1980] (a) $NH_{\perp}Cl$ The reddish brown coloured gas formed when (b) $(NH_4)_2 SO_4$ nitric oxide is oxidised by air is [IIT 1979] (c) $NH_4Cl + NaNO_3$ (d) $NH_4NO_3$ (a) $N_2O_5$ (b) $N_2O_4$ **86.** Nitrogen (I) oxide is produced by [IIT 1989] (c) $NO_2$ (d) $N_2O_3$ (a) Thermal decomposition of ammonium nitrate **98.** When $AgNO_3$ is heated strongly, the products (b) Disproportionation of $N_2O_4$ formed are (c) Thermal decomposition of ammonium nitrite [Roorkee 1990] (d) Interaction of hydroxyl amine and nitrous acid (b) $NO_2$ and $O_2$ (a) NO and $NO_2$ **87.** Which of the following is not correct for $N_2O$ [CPMT 1984] (c) $NO_2$ and $N_2O$ (d) NO and $O_2$ (a) It is called laughing gas (b) It is nitrous oxide Which of the nitrates on strong heating leaves the 99. [KCET 1990] (c) It is not a linear molecule metal as the residue (d) It is least reactive in all oxides of nitrogen (a) $AgNO_3$ (b) $Pb(NO_3)_2$

(c)  $Cu(NO_3)_2$  [NCERT 1975; AIIMS 1991] **100.** Nitrogen dioxide

(d)  $Al(NO_3)_3$ 

(a) Dissolves in water forming nitric acid

(b) Does not dissolve in water

[KCET 1989]

88. Which of the following oxides of nitrogen is the

Which of the following is a true acidic anhydride

(b)  $N_2O_3$ 

(d)  $N_2O_5$ 

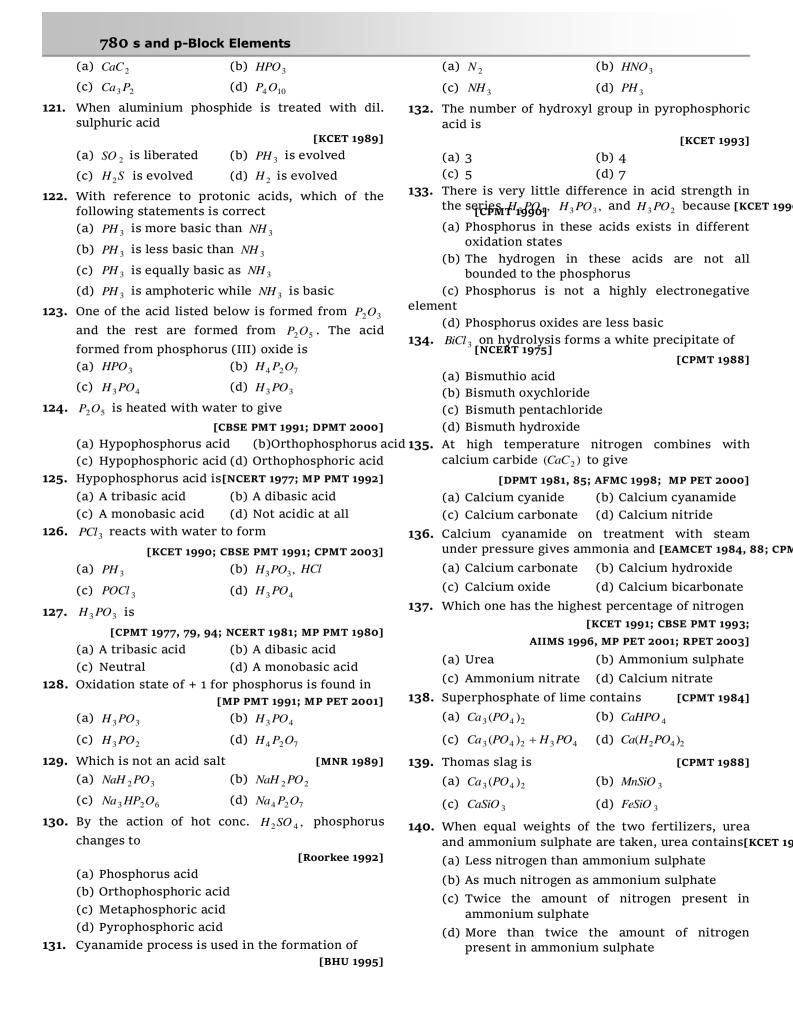
[NCERT 1977]

anhydride of nitrous acid

(a) NO

(c)  $N_2O_4$ 

(c) Dissolves in water to form nitrous acid and [KCET 1989] gives off oxygen (a) Heating it with a catalyst in an inert (d) Dissolves in water to form a mixture of atmosphere nitrous and nitric acids (b) Distilling it in an inert atmosphere 101. Concentrated nitric acid oxidises cane sugar to (c) Dissolving it in carbon disulphide [CBSE PMT 1991] crystallizing (d) Melting it and pouring the liquid into water (a)  $CO_2$  and  $H_2O$ (b) CO and  $H_2O$ 112. Bones glow in the dark because [EAMCET 1980] (c) CO,  $CO_2$  and  $H_2O$  (d) Oxalic acid and (a) They contain shining material water (b) They contain red phosphorus **102.** A mixture of ammonia and air at about  $800^{\circ} C$  in (c) White phosphorus undergoes slow combustion the presence of Pt gauze forms [Pb. CET 1989] in contact with air (b) NO (a)  $N_2O$ (d) White phosphorus changes into red form 113. Which of the following properties of white (c)  $NH_2OH$ (d)  $N_2O_3$ phosphorus are shared by red phosphorus[NCERT 1973, 74] 103. Which of the following acid possesses oxidising, (a) It shows phosphorescenes in air reducing and complex forming properties [MNR 1985] (b) It reacts with hot aqueous NaOH to give (a)  $HNO_3$ (b)  $H_2SO_4$ phosphine (c) HCl (d)  $HNO_2$ (c) It dissolves in carbon disulphide 104. Nitrogen is essential constituent of all [MP PMT 1990] (d) It burns when heated in air (a) Proteins (b) Fats 114. Mixture used for the tips of match stick is[DPMT 1984] (c) Proteins and fats (d) None of these (a) S + K105. Ammonia gas can be collected by the displacement (b)  $Sb_2S_3$ (c)  $K_2Cr_2O_7 + S + \text{red } P$ [NCERT 1989, 90] (b) Brine (d)  $K_2Cr_2O_7 + K + S$ (a) Conc.  $H_2SO_4$ 115. In modern process phosphorus is manufactured (c) Water (d) Mercury by **106.** The chemical used for cooling in refrigeration is [CPMT 1974, 78, 81] [CPMT 1981, 88] (a) Heating a mixture of phosphorite mineral (a) *CO*<sub>2</sub> (b)  $NH_4OH$ with sand and coke in electric furnace (c)  $NH_4Cl$ (d) Liquid NH3 (b) Heating calcium phosphate with coke (c) Heating bone ash with coke 107. A hydride of nitrogen which is acidic is (d) Heating the phosphate mineral with sand [NCERT 1978, 80; CPMT 1980; BHU 1986] 116. White phosphorus when boiled with strong (a)  $NH_3$ (b)  $N_2H_4$ solution of caustic soda produces [CPMT 1989,03] (c)  $N_2H_2$ (d)  $N_3H$ (a) Phosphine (b) Phosphoric acid **108.**  $PCl_5$  exists but  $NCl_5$  does not because (c) Phosphorus acid (d) No reaction [EAMCET 1977, 82] White phosphorus reacts with caustic soda. The products are  $PH_3$  and  $NaH_2PO_2$ . This reaction is (a) Nitrogen has no vacant orbitals an example of (b)  $NCl_5$  is unstable [IIT 1980; KCET 1993] (c) Nitrogen atom is much smaller (a) Oxidation (b) Reduction (d) Nitrogen is highly inert (c) Oxidation and reduction (d) Neutralisation 109. Phosphide ion has the electronic structure similar 118. Phosphine is not obtained by the reaction[MP PMT 1989] to that of (a) White P is heated with NaOH [CPMT 1988] (b) Red P is heated with NaOH (a) Nitride ion (b) Fluoride ion (c)  $Ca_3P_2$  reacts with water (c) Sodium ion (d) Chloride ion (d) Phosphorus trioxide is boiled with water 110. Which of the following phosphorus is most stable 119.  $PH_4I + NaOH$  forms [CBSE PMT 1991] [AFMC 1992] (a) Red (b) White (a)  $PH_3$ (b)  $NH_3$ (c) Black (d) All stable (c)  $P_4O_6$ (d)  $P_4 O_{10}$ 111. Red phosphorus can be obtained from white 120. Phosphine is produced by adding water to [KCET 1991] phosphorus by



- **141.** Which statement is wrong for *NO* 
  - (a) It is anhydride of nitrous acid
  - (b) Its dipole moment in 0.22 D
  - (c) It forms dimer
  - (d) it is paramagnetic
- 142. The resonance hybrid of nitrate ion is [AFMC 2002]

(a) 
$$O = N = O$$

(b) 
$$O = N = O$$
 $O = O$ 
 $O = O$ 
 $O = O$ 
 $O = O$ 

(c) 
$$O \stackrel{-1/3}{=} N \stackrel{-1/3}{=} O$$

(a) 
$$O = N = O$$
 (b)  $O = N = O$  (c)  $O = N = O$  (d)  $O = N = O$  (e)  $O = N = O$  (e)  $O = N = O$  (for  $O = O = O$  )  $O = O = O$  (for  $O = O = O$  )  $O = O = O$  (for  $O = O = O$  )  $O = O = O$  (for  $O = O = O$  )  $O = O = O$  (for  $O = O = O$  )  $O = O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O = O$  )  $O = O$  (for  $O = O$  )  $O = O$  )  $O = O$  (for  $O = O$  )  $O = O$  (for  $O = O$  )  $O = O$  )  $O = O$  (for  $O = O$  )  $O = O$  )  $O = O$  (for  $O = O$  )  $O = O$  )  $O = O$  (for  $O = O$  )  $O = O$  )  $O = O$  (for  $O = O$  )  $O = O$  )  $O = O$  (for  $O = O$  )  $O = O$  )  $O = O$  (for  $O = O$  )  $O = O$  )  $O = O$  (for  $O = O$  )  $O =$ 

- **143.** Blasting of TNT is done by mixing
  - (a) NH<sub>4</sub>Cl
- (b)  $NH_4NO_3$
- (c)  $NH_4NO_2$
- (d)  $(NH_4)_2 SO_4$
- **144.** Sodium hydroxide solution reacts with phosphorus to give phosphine. To bring about this reaction, we need

[KCET 1989]

[DPMT 2005]

- (a) White phosphorus and dil. NaOH
- (b) White phosphorus and conc. NaOH
- (c) Red phosphorus and dil. NaOH
- (d) Red phosphorus and conc. NaOH
- 145. Which of the following exhibits highest solubility in water

[MP PET 1994]

- (a)  $NH_3$
- (b)  $PH_3$
- (c)  $AsH_3$
- (d)  $SbH_3$
- 146. Which of the following has highest boiling point

[MP PET 1994]

- (a)  $NH_3$
- (b) PH<sub>3</sub>
- (c)  $AsH_3$
- (d)  $SbH_3$
- 147. In the following reaction

 $P_4 + 3NaOH + 3H_2O \rightarrow PH_3 + 3NaH_2PO_2$  [MP PMT 1994]

- (a) Phosphorus is oxidised
  - (b) Phosphorus is oxidised and reduced
  - (c) Phosphorus is reduced
  - (d) Sodium is oxidised
- **148.**  $HNO_3$  in aqueous solution yields [AMU 2000]
  - (a)  $NO_3^-$  and  $H^+$
- (b)  $NO_3^-$  and  $H_3O^+$
- (c)  $NO_2^-$  and  $OH^-$
- (d)  $N_2O_5$  and  $H_2O$
- 149. The oxyacid of phosphorus, in which phosphorus has the lowest oxidation state, is [KCET (Med.) 2001]
  - (a) Hypophosphorus acid (b)Orthophosphoric acid
  - (c) Pyrophosphoric acid (d) Metaphosphoric acid
- 150. Superphosphate is a mixture of [KCET (Med.) 2001]
  - (a)  $Ca(H_2PO_4)_2H_2O + CaCl_2.2H_2O$

- (b)  $Ca_3(PO_4)_2 . H_2O + CaCl_2 . 2H_2O$
- (c)  $Ca_3(PO_4)_2 . H_2O + 2CaSO_4 . 2H_2O$
- (d)  $Ca(H_2PO_4)_2.H_2O + 2CaSO_4.2H_2O$
- **151.** Solid  $PCl_5$  exits as

[IIPMER 2002]

- (a)  $PCl_5$
- (b) *PCl*<sub>4</sub><sup>+</sup>
- (c) *PCl*<sub>6</sub>
- (d)  $PCl_4^+$  and  $PCl_6^-$
- **152.** In the reaction,  $P_2O_5 + 3CaO \rightarrow Ca_3(PO_4)_2; P_2O_5$ acts as..... [Orissa JEE 2002]

(a) Acidic flux

- (b) Basic flux
- (c) Basic impurity
- (d) Acidic impurity
- **153.** Atoms in a  $P_4$  molecule of white phosphorus are arranged regularly in space in which of the following way

[Kerala (Engg.) 2002]

- (a) At the corners of tetrahedron
- (b) At the corners of a cube
- (c) At the corners of a four membered ring
- (d) At the centre and corners of an equilateral triangle
- 154. The most common minerals of phosphorus are

[Kerala (Med.) 2002]

- (a) Hydroxy apatite and kernite
- (b) Colemanite and fluoraptite
- (c) Borax and fluoraptite
- (d) Hydroxy apatite and colemanite
- (e) Hydroxy apatite and fluorapatile
- **155.** The three important oxidation states of phosphorus are

[Kerala (Med.) 2002]

- (a) -3.+3 and +5
- (b) -3, +3 and -5
- (c) -3.+4 and -4
- (d) -3.+3 and +4
- **156.** In case of nitrogen,  $NCl_3$  is possible but not  $NCl_5$ while in case of phosphorous, PCl3 as well as PCl<sub>5</sub> are possible. It is due to
  - (a) Availability of vacant d-orbital in P but not in
  - (b) Lower electronegativity of P than N
  - (c) Lower tendency of *H* bond formation in *P* than
  - (d) Occurrence of *P* in solid while *N* in gaseous state at room temperature
- 157. When ammonia is passed over heated copper oxide, the metallic copper is obtained. reaction shows that ammonia is
  - (a) A dehydrating agent (b) An oxidising agent
  - (c) A reducing agent
    - (d) A nitrating agent
  - Urea is preferred to ammonium sulphate as a nitrogeneous fertilizer because
    - (a) It is more soluble in water
    - (b) It is cheaper than ammonium sulphate

- (c) It is quite stable
- (d) It does not cause acidity in the soil
- 159. Liquid ammonia is used for refrigeration because

#### [MP PET 2002]

- (a) It has a high dipole moment
- (b) It has a high heat of vapourisation
- (c) It is basic
- (d) It is a stable compound
- **160.** Action of concentrated nitric acid (HNO<sub>3</sub>) on metallic tin produces [BHU 2002]
  - (a) Stannic nitrate
- (b) Stannous nitrate
- (c) Stannous nitrite
- (d) Meta stannic acid
- 161. How can you synthesize nitric oxide in the laboratory

#### [Orissa JEE 2003]

- (a) Zinc with cold and dilute HNO<sub>2</sub>
- (b) Zinc with concentrated HNO<sub>3</sub>
- (c) Copper with cold and dilute HNO<sub>3</sub>
- (d) Heating  $NH_4NO_3$
- 162. What would happen when a solution of potassium chromate is treated with an excess of dilute nitric [AIEEE 2003]
  - (a)  $Cr^{3+}$  and  $Cr_2O_7^{2-}$  are formed
  - (b)  $Cr_2O_7^{2-}$  and  $H_2O$  are formed
  - (c)  $CrO_4^{2-}$  is reduced to +3 state of Cr
  - (d)  $CrO_4^{2-}$  is oxidized to +7 state of Cr
- **163.** The pentavalence in phosphorus is more stable as compared to that of nitrogen even though they belong to the same group. It is due to
  - (a) Inert nature of nitrogen
  - (b) Reactivity of phosphorus
  - (c) Larger size of phosphorus atom
  - (d) Dissimilar electronic configuration
- **164.** A neutral fertilizer among the following compounds is

[KCET 2002]

- (a) Urea
- (b) Ammonium nitrate
- (c) Ammonium sulphate
- (d) Calcium ammonium nitrate
- 165. Which is true with regard to the properties of

[BHU 2000]

- (a)  $PH_3$  is not much stable
- (b)  $PH_3$  is neutral towards litmus
- (c)  $PH_3$  has fishy smell
- (d)  $PH_3$  is insoluble in water
- **166.** Nitrogen is obtained when NaNO, reacts with

#### [UPSEAT 2003]

- (a)  $NH_{\perp}Cl$
- (b)  $NH_4NO_3$
- (c)  $(NH_4)_2CO_3$
- (d)  $NH_4OH$

- **167.**  $N_2$  combines with metal to form [JIPMER 2000]
  - (a) Nitride
- (b) Nitrate
- (c) Nitrite
- (d) Nitrosyl chloride
- **168.** The number of P-O-P bridges in the structure of phosphorous pentoxide and phosphorous trioxide are respectively [AIIMS 2005]
  - (a) 6, 6
- (b) 5, 5
- (c) 5, 6
- (d) 6, 5
- 169. Sodium pyrophosphate is
  - (b)  $Na_{4}P_{2}O_{7}$
  - (a)  $Na_2P_2O_7$
- (d)  $Na_2PO_2$
- (c)  $NaPO_{A}$
- 170. Which of the following is solid in nature

#### [UPSEAT 2003; AFMC 2004]

- (a) NO
- (b)  $N_2O$
- (c)  $N_2O_3$
- (d)  $N_2O_5$
- 171. Which of the following is a cyclic phosphate

[CPMT 2003]

- (a)  $H_5P_3O_{10}$
- (b)  $H_6 P_4 O_{13}$
- (c)  $H_5P_5O_{15}$
- (d)  $H_7 P_5 O_{16}$
- 172. The reaction, which forms nitric oxide, is

#### [KCET (Med.) 2001]

- (a) C and  $N_2O$
- (b) Cu and  $N_2O$
- (c) Na and  $NH_3$
- (d) Cu and  $HNO_3$
- 173. Which one of the following can be used as an anaesthetic

#### [EAMCET 1998]

- (a)  $N_2O$
- (b) NO
- (c)  $NCl_3$
- (d)  $NO_2$
- 174. Solution of sodium metal in liquid ammonia is strongly reducing due to the presence in the solution of the following

#### [KCET 2000; MP PMT 2001]

- (a) Sodium hydride
- (b) Sodium amide
- (c) Sodium atoms
- (d) Solvated electrons
- 175. What may be expected to happen when phosphine gas is mixed with chlorine gas [AIEEE 2003]
  - (a) The mixture only cools down
  - (b) PCl3 and HCl are formed and the mixture warms up
  - (c) PCl<sub>5</sub> and HCl are formed and the mixture cools down
  - (d)  $PH_3.Cl_2$  is formed with warming up
- **176.**  $P_4O_{10}$  is not used with to dry  $NH_3$  gas because

#### [KCET 2001]

- (a)  $P_4O_{10}$  is basic and  $NH_3$  is acidic
- (b)  $P_4O_{10}$  is acidic and  $NH_3$  is basic
- (c)  $P_4O_{10}$  is not a drying agent
- (d)  $P_4O_{10}$  reacts with moisture in  $NH_3$

177.		with sodium hypochlorite,	189.		compounds is sparingly
	product containing nitro (a) $N_2$	_		soluble in ammonia	[JIPMER 1999]
	=	(b) $N_2O$		(a) $AgI$	(b) $AgBr$
	=			(c) AgCl	(d) CuCl <sub>2</sub>
178.	$P_2O_5$ is used extensively (a) Reducing agent		190.	The carbonate which do heating is	es not leave a residue on
	(c) Dehydrating agent				[JIPMER 1999; DCE 1999]
179.		due to [DCE 2000; MP PET 2001	1	(a) $Na_2CO_3$	(b) $Ag_2CO_3$
, ,	(a) No vacant <i>d</i> -orbital	,	-	(c) $CuCO_3$	(d) $(NH_4)_2 CO_3$
	(b) High dissociation en (c) High electronegativi		191.	Which of the following moment	g has the highest dipole
	(d) None	,			[CBSE PMT 1997]
180.	Which show maximum v	valency [CPMT 2003]		(a) $NH_3$	(b) $PH_3$
	(a) Phosphorus	(b) Tin		(c) $SbH_3$	(d) $AsH_3$
	(c) Antimony	(d) Bismuth	192.	The structural formula o	f hypophosphorus acid is
181.	Which is used in the manufacture of $NH_3$	Haber process for the [MH CET 2001]			BCECE 2005; Pb. CET 2002]
	_				
	(a) Pt (c) CuO	(b) $Fe + Mo$ (d) $Al_2O_3$		(a) $H \nearrow P$ OH	(b) $P OH$
	• •	2 3		H	OH
182.		mmonium hydroxide to a	Γ'n	(a) $H \stackrel{\mid}{\stackrel{\mid}{\stackrel{\mid}{\stackrel{\mid}{\stackrel{\mid}{\stackrel{\mid}{\stackrel{\mid}{\stackrel{\mid}$	o
	copper chloride solution (a) A deep blue solution		ΓN	(c) < R	(d) / P
	(b) No change is observe			$H \cap OH$	$OH \longrightarrow OOH$
	•	of copper hydroxide is		11	OII
	obtained	or copper hydroxide is	193.	-	of the following fertilizers
	(d) Black precipitate of c	opper oxide is obtained			ty of the soil[CBSE PMT 1998]
183.		by heating $(NH_4)_2 SO_4$ and		(a) Urea	(b) Potassium nitrate
	KCNO is		lime	(c) Ammonium sulphate	(d) Superphosphate of
		[DPMT 2000]		Aqua-regia is	[Orissa JEE 2005]
	(a) Hydrocyanic acid	(b) Ammonia	134.	(a) 1:3 conc. $HNO_3$ and	
	(c) Ammonium cyanide			(b) 1: 2 conc. $HNO_3$ and	
184.		O-P bonds in cyclic		-	
		[IIT-JEE (Screening) 2000]		(c) $3:1$ conc. $HNO_3$ and	
	<ul><li>(a) Zero</li><li>(c) Three</li></ul>	(b) Two		(d) 2:1 conc. $HNO_3$ and	conc. HCl
10-		(d) Four	195.	_	n is obtained on heating
105.	*	into the palm and washed		ammonium nitrate at 250	0 ° C
	presence of	nto yellow. It shows the		(a) Nitric oxide	(b) Nitrous oxide
	presence of	[CPMT 1997]		(c) Nitrogen dioxide	(d) Dinitrogen oxide
	(a) $NO_2$	(b) $N_2O$	196.	The oxidation number of	
	(c) <i>NO</i>	(d) $N_2O_5$			shetra CEE 1998; DCE 2001]
106				(a) -1 to +3	(b) -3 to +3
100.	Which of the following i	[CPMT 1999]	10=	(c) -3 to +5	(d) -5 to +1
	(a) Bone meal	(b) Thomas meal	197.	Inorganic graphite is	[KCET 2003]
	(c) Nitro phosphate	(d) Ammonium sulphate		(a) $B_3 N_3 H_6$	(b) $B_3N_3$
187	Which compound is rela	<u>-</u>		(c) SiC	(d) $Fe(CO)_5$
10/.	vvincii compound is reid	[RPET 1999]	198.	Conc. $HNO_3$ can be store	d in container of[MH CET 2001]
	(a) <i>CO</i> <sub>2</sub>	(b) $H_2$		(a) Al	(b) <i>Sn</i>
	(c) $NO_2$			(c) Cu	(d) <i>Zn</i>
-00	=	(d) NH <sub>3</sub>	199.	Which of the following c	_
188.		[CPMT 2002; JIPMER 2002]		(a) SLCI	[JIPMER 1997]
	(a) Quick lime	(b) Slaked lime		(a) SbCl <sub>3</sub>	(b) <i>BiCl</i> <sub>5</sub>
	(c) Anhy. $CaCl_2$	(d) None of these		(c) SbCl <sub>5</sub>	(d) $AsCl_5$

200.	The formula of an $H_3PO_4$ . It is a	n oxyacid of phosphorus is	212.	One mole of calcium phosphide on reaction with excess water gives [IIT-JEE 1999]
	[MP PMT 19	96; CPMT 1999; J & K CET 2005]		(a) One mole of phosphine
	(a) Dibasic acid	(b) Monobasic acid		(b) Two moles of phosphoric acid
	(c) Tribasic acid	(d) Tetrabasic acid		(c) Two moles of phosphine
201.	Which salt can be cl	assified as an acid salt[ <b>CPMT</b> 19	93]	(d) One mole of phosphorous pentoxide
	(a) $Na_2SO_4$	(b) BiOCl	213.	On heating ammonium dichromate, the gas
	(c) Pb(OH)Cl	(d) $Na_2HPO_4$		evolved is
202.	In $NH_3$ and $PH_3$ , th	<del>-</del> · · ·		[IIT-JEE 1999] (a) Oxygen (b) Ammonia
	(a) Odour	(b) Combustibility		(c) Nitrous oxide (d) Nitrogen
	(c) Basic nature	(d) None of these	214	In compounds of type $ECl_3$ , where $E = B$ , $P$ , $As$ or
203.		following hydrides is least		Bi, the angles $Cl - E - Cl$ for different $E$ are in the
		(b) ShH		order
	(a) $AsH_3$	(b) SbH <sub>3</sub>		[HT-JEE 1999]
	(c) $NH_3$	(d) $PH_3$		(a) $B > P = As = Bi$ (b) $B > P > As > Bi$
204.		m V group, gives most basic		(c) $B < P = As = Bi$ (d) $B < P < As < Bi$
	compound with hydr	_	215[1	տ <b>Aբարթդiց</b> 59n reaction with hypochlorite anion, can form
	(a) Nitrogen	(b) Bismuth		[IIT-JEE 1999]
	(c) Arsenic	(d) Phosphorus		(a) NO (b) $NH_4Cl$
205.		is two series of salts is[KCET 19	96]	(c) $N_2H_4$ (d) $HNO_2$
	(a) $H_3PO_4$	(b) $H_3PO_3$	216	Orthophosphoric acid represents the molaysis
	(c) $H_3BO_3$	(d) $H_3PO_2$	210.	condition due to
206.	6. Which gas is obtained when urea is heated with			(a) Hydrogen bonding
	$HNO_2$			(b) Phosphorous group
		[CPMT 1996]		(c) Maximum oxygen group
	(a) $N_2$	(b) $H_2$		(d) Tribasicity
	(c) $O_2$	(d) $NH_3$	217.	Which of the following elements forms a strongly
207.	Atomic number of A	V is 7. The atomic number of		acidic oxide
	IIIrd member of nitr	rogen family is [CPMT 1996]		(a) P (b) As (c) Sb (d) Bi
	(a) 23	(b) 15	218	In group V-A of the periodic table nitrogen forms
	(c) 33	(d) 43	210.	only a trihalide but other elements form
208.		ring have least covalent $P-H$		pentahalides also. The reason is
	bond	[CDWT 1006]		(a) Nitrogen has less affinity towards halogens
	(a) $PH_3$	[CPMT 1996] (b) $P_2H_6$		(b) Nitrogen halides are covalent
	-			(c) Nitrogen undergoes $d^2sp^3$ hybridization
	(c) $P_2H_5$	(d) $PH_6^+$		(d) Nitrogen does not have <i>d</i> -orbitals
209.	Sides of match box h (a) Potassium chlora	_	219.	In the nitrogen family the $H-M-H$ bond angle in the hydrides $MH_3$ gradually becomes closer to
		ate, antimony sulphide		90° on going from $N$ to $Sb$ . This shows that
	(c) Antimony sulphi			gradually
	(d) Antimony sulphi	de, red lead		[MP PET/PMT 1998; MP PMT 2000]
210.	Which of the followi	ng compound is tribasic acid		(a) The basic strength of hydrides increases
		[AIIMS 2002]		(b) Almost pure $p$ -orbitals are used for $M-H$
	(a) $H_3PO_2$	(b) $H_3PO_3$	bond	
	(c) $H_3PO_4$	(d) $H_4 P_2 O_7$		(c) The bond energies of $M-H$ bond increase (d) The bond pairs of electrons become nearer to
211.	Which of the follow molecular nitrogen l	ring is manufactured from the by bacteria [MP PET 1999]		the central atom
	(a) $NO_3$	(b) $NO_2$	220.	An element (X) forms compounds of the formula $XCl_3$ , $X_2O_5$ and $Ca_3X_2$ , but does not form $XCl_5$ ,
	(c) Amino acids	(d) Ammonia		which of the following is the element $X$ [MP PET 1997]
	(5) Immilio deldo			(a) $B$ (b) $Al$

					s a	nd p-Block	Elements 7	785
	(c) N	(d) P			(a) N		(b) P	
221.	Which of the follounchanged on going				(c) As		(d) <i>Sb</i>	
	(Group-VA)?	Jown III the II	itti ogen Tanniy	233.	The boiling p	oints of the	e following h	ydrides follow
	(010 up 111) 1		[MP PMT 1997]		the order of			[DPMT 2004]
	(a) Highest oxidation	state			(a) $NH_3 > AsH$	H. > PH. > S	hН .	[DFM1 2004]
	(b) Non-metallic char				-		-	
	(c) Stability of hydrid	les			(b) $SbH_3 > As$	-	-	
	(d) Physical state		h h		(c) $SbH_3 > NI$	5	3	
222.	Which of the followin a reducing agent and		phosphorus is	Г1	(d) $NH_{3} > PH$ <b>DCE 2004</b> ]	$S_3 > AsH_3 > S$	$bH_3$	
	(a) $H_3PO_2$	(b) $H_3PO_3$		234.	Which of the	following l	nalides is mos	st acidic
	(c) $H_3PO_4$	(d) $H_4 P_2 O_6$					_	MP PMT 2004]
222	Bone black is a polym				(a) $PCl_3$		(b) $BiCl_3$	
223.	(a) Phosphorus	(b) Sulphu			(c) $SbCl_3$		(d) <i>CCl</i> <sub>4</sub>	
	(c) Carbon	(d) Nitrog		235.	In the elect	trothermal	process, th	ne compound
224.	Nitrous oxide is know		[AFMC 2004]		displaced by	silica from	calcium phos	sphate is [KCET
	(a) Breathing gas	(b) Laughi			(a) Calcium		(b) Phosphi	ne
		(d) Labora			(c) Phosphor	us	(d) Phospho	orus
225.	When lead nitrate is h				oxide			
	(a) $NO_2$	(b) <i>NO</i>		236.	Which of sublimation	the follo	wing comp	ound show
	(c) $N_2O_5$	(d) $N_2O$			Subililiation		[AEMC 1005.	Pb. CET 2000]
226	Which element exist	-	at 25°C and 1		(a) NH <sub>4</sub> Cl		(b) $CaCO_3$	FU. CET 2000]
220.	atmospheric pressure			03]			,	
	(a) <i>Br</i>	(b) <i>Cl</i>	8 2	0.	(c) $BaSO_4$		(d) $CaHPO_3$	
	(c) Hg	(d) P		237.	Number of P	− <i>O</i> bonds i	n $P_4O_{10}$ is	[DCE 2002]
227.	In the reaction H	$NO_3 + P_4O_{10} \rightarrow 4$	$HPO_3 + x$ , the		(a) 17		(b) 16	
	product x is				(c) 15		(d) 6	
		[MH CET 200	03; DPMT 2004]	238.	Most acidic o	xide is	I	Pb. CET 2004]
	(a) $N_2O_3$	(b) $N_2O_5$			(a) $Na_2O$		(b) <i>ZnO</i>	
	(c) $NO_2$	(d) $H_2O$			(c) MgO		(d) $P_2O_5$	
228.	The number of hydrophosphorus atom in h				Which of the	following i		hydride [ <b>Pb. CET 2003]</b>
	(a) Zero	(b) Two			(a) <i>NO</i>		(b) <i>NO</i> <sub>2</sub>	
	(c) One	(d) Three					(d) $N_2O$	
229.	Which blue liquid				(c) $N_2O_5$		-	
	equimolar amounts of		–30°C [IIT-JEE (S	cr <b>eq10</b> 1	n <b>g)xidasil</b> on nu	mber of As	in $H_2 AsO_4^-$ is	[CPMT 2001]
	(a) $N_2O$	(b) $N_2O_3$			(a) 6		(b) 7	
	(c) $N_2O_4$	(d) $N_2O_5$			(c) 5		(d) 9	
230.	Which is the most		•		When plants			_
	allotropic form of pho	-	EE (Screening) 200	05]	-		_	nitrogen. The
	(a) Red	(b) White			inorganic nit			[KCET 2005]
	(c) Black	(d) Yellow			(a) Ammonia			s of nitrogen
231.	$(NH_4)_2 Cr_2 O_7$ on heating				(c) Nitrates		(d) Nitrides	<b>;</b>
	gas will be obtained by (a) Heating $NH_4NO_2$	y [IIT JEE (Scr	eening) 2004; BVP	2004]		Oxygen	family	
	(b) Heating $NH_4NO_3$							
	(c) Treating $H_2O_2$ wire	th <i>NaNO</i> <sub>2</sub>		1.	Which elemen	nt is found	in free state	
		_				[CP]	MT 1972, 81, 9	1; DPMT 1986]
	(d) Treating $Mg_3N_2$ w	_	11 11 11		(a) Iodine		(b) Sulphur	
232.	The element which for states $+I$ to $+V$ is	orms oxides i	n all oxidation [AIIMS 2004]		(c) Phosphor	us	(d) Magnes	ium

	, 00 5 and p 2100.						
2.	Which of the element allotropic forms	s listed below	occurs in		(a) H <sub>2</sub> O	(b) H <sub>2</sub> S	
	anotropic forms	[C	PMT 1972]		(c) $H_2Se$	(d) $H_2Te$	
	(a) Iodine	(b) Copper	1 19/2]	16.	Oxygen was discover	-	[BHU 1987]
	(c) Sulphur	(d) Silver			(a) Priestley	(b) Boyle	
2	•		EMC 100=1		(c) Scheele	(d) Cavano	
3.	Which forms new comp		FMC 1987]	17.	The compound which	n gives off oxyge	en on moderate
	(a) $H_2O$ in air	(b) $O_2$ in air			heating is	MP PMT 1991; M	ADT Ribar 100El
	(c) $N_2$ in air	(d) Phosphorus	in air		(a) Cupric oxide	WIF FWII 1991, WI	ADI Biliai 1995]
4.	Which of the following	g after burning	at room		(b) Mercuric oxide		
	temperature gives gase	ous oxide			(c) Zinc oxide		
	[NCERT 1973; CPMT 198	1; DPMT 1982; JIP	MER 2001]		(d) Aluminium oxide	9	
	(a) <i>H</i>	(b) <i>Na</i>		18.	It is possible to		from air by
	(c) S	(d) <i>He</i>			fractional distillation		•
5.	Sulphur molecule is co		phur ion.		(a) Oxygen is in a d	ifferent group	of the periodic
<b>J</b> .	when it		,		table from nitrog		
		[D:	PMT 1980]		(b) Oxygen is more		
	(a) Gains two electrons				(c) Oxygen has high		
	(c) Gains two protons				(d) Oxygen has a lov	-	_
6.	The element which li			19.	Oxygen is denser tha		
•	water is	our acco only gen	840 110111		(a) II O		o; MP PET 1999]
		[MP	PMT 1993]		(a) $H_2O$	(b) Ethano	
	(a) <i>P</i>	(b) <i>Na</i>			(c) Mercury	(d) Kerose	ne oil
	(c) F	(d) <i>I</i>		20.	Oxygen molecule exl		
7•	The highest catenation	-	-			9, 2002; AIIMS 1 1; MP PMT 1985,	
		3; MP PET 1993; C	PMT 1997]		(a) Paramagnetism		
	(a) Oxygen	(b) Sulphur			(c) Ferromagnetism		-
	(c) Selenium	(d) Tellurium		21.	When oxygen is p		_
8.	Ozone belongs to whitable		e periodic		$Na_2SO_3$ we get		
	(a) V group	(b) VI group			(a) $Na_2SO_4$	(b) $Na_2S$	
	(c) VII group	(d) None of the			(c) $NaHSO_4$	(d) <i>NaH</i>	
9.	The number of unpa		n the <i>p</i> -	22.	Oxygen does not rea K <b>urukshetra CEE 1991]</b> (a)	ct with [	CBSE PMT 1989]
	subshell of oxygen aton			LH	(a) Petra CEE 1991]	(b) <i>Na</i>	
	(a) 1	(b) 2 (d) 4			(c) S	(d) <i>Cl</i>	
10.	(c) 3 Most abundant element		ET/DMT 1000	23.	The formula of ozon	e is $O_3$ , it is	
10.	(a) O	(b) $S$	E1/FWI 1900]	J	[CPMT 1989, 91; Mani]	pal MEE 1995; R	PET 1999, 2000]
	(c) Se	(d) <i>Te</i>			(a) An allotrope of o	xygen (b)Con	npound of oxygen
11.	Which of the following		gative[BHU 10	9781	(c) Isotope of oxyge		of these
	(a) <i>O</i>	(b) S	8	24.	Ozone is obtained fr		[CPMT 1982]
	(c) <i>Te</i>	(d) <i>Se</i>			(a) By oxidation at h	-	re
12.	The ability of a substar		o or more		(b) By oxidation using		
	crystalline structures is				(c) By silent electric	discharge	
	(a) Isomerism	(b) Polymorphi	sm		(d) By conversion at		ia tura aleant
	(c) Isomorphism	(d) Amorphism		25.	Which of the follow ozone layer	wing statement	is true about
13.	$SO_2$ is obtained when	[Roo	rkee 1995]		Ozone layer		[NCERT 1980]
	<ul><li>(a) Oxygen reacts with</li><li>(b) Hydrolysis of dilute</li></ul>	=	acid		(a) It is harmful be	ecause ozone is	
	(c) Concentrated $H_2SO$	= .	$_2SO_3$		(b) It is beneficial b		
	(d) All of these				proceed faster in	_	
14.	Which shows polymorp	hism <b>[BHU 1982; M</b>	IP PMT 1985]		(c) It is beneficial ultraviolet radia		cuts out the
	(a) <i>O</i>	(b) S			uiti avioiet radla	tion of the sun	
	(c) <i>Se</i>	(d) All the abov	re				
15.	Bond angle is minimum	for [D	PMT 1990]				

		ause ozone cuts out the n of the sun which are vital	36.	5. A solution of sulphur dioxide in water reacts with $H_2S$ precipitating sulphur. Here sulphur dioxide acts as			
26.		statement with respect to			[NCERT 1980; I	// P PMT 100/1	
	ozone			(a) An oxidising agent			
		[AIIMS 1992]		(c) An acid	(d) A reducin		
		the upper atmosphere by a	37.	When $SO_2$ is passed t	-		
	photochemical reac	tion involving dioxygen	3/•	solution	in ough dordin	100 H <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub>	
	(b) Ozone is more react	tive than oxygen		Solution	[CI	PMT 1989, 94]	
		etic whereas dioxygen is		(a) The solution turns b		WI 1909, 94]	
	paramagnetic			(b) The solution is deco			
		ne earth's inhabitants by		(c) $SO_2$ is reduced	iourisea		
	absorbing $\gamma$ radiat			=	·		
27.		ving property is not correct		(d) Green $Cr_2(SO_4)_3$ is f			
	for ozone	[CD25m + -0 +1	38.	When $SO_2$ is passed	through cup	ric chloride	
	(-) The sector and sector	[CPMT 1984]		solution			
	(a) It oxidises lead sulp					79, 81, 89, 94]	
	(b) It oxidises potassiu			(a) A white precipitate			
	(c) It oxidises mercury			(b) The solution becomes colourless			
	(d) It cannot act as blea			(c) The solution becomes colourless and a white			
28.	Ozone with $KI$ solution	-		precipitate of $Cu_2Cl$			
	(a) <i>Cl</i> <sub>2</sub>	(b) $I_2$		(d) No visible change ta	_		
	(c) HI	(d) $IO_3$	39.	Which of the following i	s oxidised by	SO <sub>2</sub> [BHU 1986]	
29.		ly absorbed by alkaline		(a) <i>Mg</i>	(b) $K_2 C r_2 O_7$		
		nnamon is [CBSE PMT 1989]		(c) $KMnO_4$	(d) All of the	ese	
	(a) $O_{3}$ , $CH_{4}$	(b) $O_2$ , $O_3$	40.	Bleaching action of $SO_2$			
	(c) $SO_2$ , $CH_4$	(d) $N_2O$ , $O_3$	40.	_			
30.	Ozone turns trimethyl j			(a) Reduction	(b) Oxidation		
,	(a) Green	(b) Violet		(c) Hydrolysis	(d) Its acidio		
	(c) Red	(d) Black	41.	A salt of sulphurous acid		[NCERT 1978]	
31.				(a) Sulphate	(b) Sulphura	te	
, 1.	No. of atoms in one molecule of sulphur is [AFMC 1987, 91; AMU 1985]			(c) Sulphite	(d) Sulphide		
	(a) 8	(b) 4	42.	Which of the following i			
	(c) 3	(d) None of these		(2) 50	[AFMC 1990;	JIPMEK 1997]	
32.		through acidified $KMnO_4$ ,		(a) <i>SO</i> <sub>3</sub>	(b) $N_2O$		
	we get	4,		(c) <i>BeO</i>	(d) <i>HgO</i>		
	_	CPMT 1979, 91; MP PMT 1987]	43.	The final acid obtained		nufacture of	
	(a) $K_2SO_3$	(b) $MnO_2$		$H_2SO_4$ by contact proce	ess is		
	(c) $KHSO_3$	(d) Sulphur		(a) $H_2SO_4$ (conc.)	(b) $H_2SO_4$ (	dil.)	
	-	-		(c) $H_2SO_4$	(d) $H_2S_2O_7$		
33.	copper turnings when	heated with concentrated [AFMC 1987; BHU 1999; CBSE I	ом <b>4</b> 4-			[CPMT 1985]	
		(b) SO <sub>3</sub>	-WI F 200	(a) Reducing agent	(b) Dehydra		
	(a) SO <sub>2</sub>	9		(c) Sulphonating agent	-		
	(c) $H_2S$	(d) $O_2$	45.	In the reaction	(4) 11161119 11		
34.	Which of the following dioxide	g is used to absorb sulphur	40.	$2Ag + 2H_2SO_4 \rightarrow Ag_2SO_4$	$+2H_2O+SO_2$		
		[EAMCET 1980]		$H_2SO_4$ acts as		[CPMT 1981]	
	(a) Conc. $H_2SO_4$	(b) KOH solution		(a) Reducing agent	(b) Oxidising		
	(c) Water	(d) Anhydrous CaCl <sub>2</sub>	_	(c) Catalytic agent	(d) Dehydra		
35.		as an oxidising as well as	46.	In the		reaction	
٠.,	reducing agent	[IIT 1991]		$HCOOH \xrightarrow{H_2SO_4} CO + H$	$U_2O$ ; $H_2SO_4$ act	s as	
	(a) $SO_2$	(b) $MnO_2$		(a) Dehydrating agent	(b) Oxidising	g agent	
	(c) $Al_2O_3$	(d) $CrO_3$		(c) Reducing agent	(d) All of the	ese	
	$(0)$ $n_2 o_3$	(a) C/O <sub>3</sub>					

<del>4</del> 7•	When conc. $H_2SO_4$ comes in contact with sugar,
	it becomes black due to
	F67777

#### [CPMT 1989; BHU 1986; MDAT Bihar 1980]

- (a) Hydrolysis
- (b) Hydration
- (c) Decolourisation
- (d) Dehydration
- **48.** Oxalic acid when heated with conc.  $H_2SO_4$ , gives out

#### [DPMT 1981; AFMC 1998]

- (a)  $H_2O$  and  $CO_2$
- (b) CO and  $CO_2$
- (c) Oxalic sulphate
- (d)  $CO_2$  and  $H_2S$
- 49. Which one is known as oil of vitriol

#### [CPMT 1988; MP PMT 2004]

- (a)  $H_2SO_3$
- (b)  $H_2SO_4$
- (c)  $H_2S_2O_7$
- (d)  $H_2S_2O_8$
- 50. The acid used in lead storage cells is

#### [NCERT 1971; Roorkee 1989]

- (a) Phosphoric acid
- (b) Nitric acid
- (c) Sulphuric acid
- (d) Hydrochloric acid
- **51.** Which one of the gas dissolves in  $H_2SO_4$  to give oleum

#### [CPMT 1988]

- (a)  $SO_2$
- (b)  $H_2S$
- (c)  $S_2O$
- (d)  $SO_3$
- **52.** Oleum is[CBSE PMT 1991; MP PMT 2002; CPMT 2004]
  - (a) Castor oil
- (b) Oil of vitriol
- (c) Fuming  $H_2SO_4$
- (d) None of them
- **53.** There is no S S bond in

#### [IIT 1991; CPMT 1999; DCE 2000]

- (a)  $S_2 O_4^{2-}$
- (b)  $S_2O_5^{2-}$
- (c)  $S_2O_3^{2-}$
- (d)  $S_2 O_7^{2-}$
- **54.** Which of the following sulphate is insoluble in water

#### [MP PMT 2000]

- (a)  $CuSO_4$
- (b)  $CdSO_4$
- (c)  $PbSO_{A}$
- (d)  $Bi_2(SO_4)_3$
- 55. When sulphur is boiled with  $Na_2SO_3$  solution, the compound formed is [CPMT 1979; Roorkee 1992]
  - (a) Sodium sulphide
- (b) Sodium sulphate
- (c) Sodium persulphate (d) Sodium thiosulphate
- **56.** The products of the chemical reaction between  $Na_2S_2O_3$ ,  $Cl_2$  and  $H_2O$  are **[EAMCET 1989]** 
  - (a)  $S + HCl + Na_2S$
- (b)  $S + HCl + Na_2SO_4$
- (c)  $S + HCl + Na_2SO_3$
- (d)  $S + NaClO_3 + H_2O$
- **57.** Hypo is used in photography for **[Roorkee 1989]** 
  - (a) Developing a picture
  - (b) Picture printing
  - (c) The colour of picture
  - (d) The fixation of the picture
- 58. Hypo is used in photography to

#### [CBSE PMT 1988; Pb. CET 1989]

(a) Reduce AgBr grains to metallic silver

- (b) Convert the metallic silver to silver salt
- (c) Remove undecomposed silver bromide as a soluble complex
- (d) Remove reduced silver
- 59. Hypo is used in photography because of its
  [IIT 1981; EAMCET 1988; MADT Bihar 1995]
  - (a) Reducing behaviour
  - (b) Oxidising behaviour
  - (c) Complex forming behaviour
  - (d) Reaction with light
- **60.** Aqueous solutions of hydrogen sulphide and sulphur dioxide when mixed together, yield[KCET 2002]
  - (a) Sulphur and water
  - (b) Sulphur trioxide and water
  - (c) Hydrogen peroxide and sulphur
  - (d) Hydrogen and sulphurous acid
- **61.** An example of a neutral oxide is **[KCET 2003]** 
  - (a) NO
- (b) CO<sub>2</sub>
- (c) CaO
- (d) ZnO
- **52.** Which of the following is the best scientific method to test presence of water in a liquid [AIIMS 1999]
  - (a) Taste
  - (b) Smell
  - (c) Use of litmus paper
  - (d) Use of anhydrous copper sulphate
- **63.**  $H_2S$  react with  $O_2$  to form [AFMC 1995]
  - (a)  $H_2O + S$
- (b)  $H_2O + SO_2$
- (c)  $H_2O + SO_3$
- (d)  $H_2SO_4 + S$
- **64.** Which of the following mixture is chromic acid **[Pb. PMT 2000]** 
  - (a)  $K_2Cr_2O_7$  and conc.  $H_2SO_4$
  - (b)  $K_2Cr_2O_7$  and HCl
  - (c)  $K_2SO_4$  and conc.  $H_2SO_4$
  - (d)  $H_2SO_4$  and HCl
- **65.** At room temperature  $H_2O$  is a liquid while  $H_2S$  is a gas. The reason is **[RPET 1999]** 
  - (a) Electronegativity of O is greater than S
  - (b) Difference in the bond angles of both the molecules
  - (c) Association takes place in  $H_2O$  due to  $H_2O$  bonding while no  $H_2O$  bonding in  $H_2O$
  - (d) O and S belong to different periods
- **66.** Which of the following products is formed on boiling tin with an alkali solution [Roorkee 2000]
  - (a)  $Sn(OH)_2$
- (b)  $Sn(OH)_4$
- (c)  $SnO_3^{2-}$
- (d)  $SnO_2$
- **67.** Amongst  $H_2O$ ,  $H_2S$ ,  $H_2Se$  and  $H_2Te$  the one with the highest boiling point is[IIT-JEE (Screening) 2000]
  - (a)  $H_2O$  because of hydrogen bonding
  - (b)  $H_2$  Te because of higher molecular weight
  - (c)  $H_2S$  because of hydrogen bonding

- (d)  $H_2Se$  because of lower molecular weight
- **68.** Among the hydrides formed by the group VI-A elements, only  $H_2O$  has an abnormally low volatility (high boiling point). This is so because
  - (a)  $H_2O$  molecules are associated due to intermolecular hydrogen bonds
  - (b)  $H_2O$  is covalent in nature
  - (c) The O-H bond in  $H_2O$  is very strong
  - (d) The electronegativity difference of  ${\it H}$  and  ${\it O}$  is very large
- **69.** Sulphuric acid reacts with *PCl*<sub>5</sub> to give

[KCET 1996; JIPMER 2000]

- (a) Thionyl chloride monochloride
- (b) Sulphur
- (c) Sulphuryl chloride
- (d) Sulphur tetrachloride
- **70.** Carbogen is

[EAMCET 1998]

- (a) Pure form of carbon
- (b) COCl<sub>2</sub>
- (c) Mixture of CO and CO,
- (d) Mixture of  $O_2$  and  $CO_2$
- **71.** Which of the following dissociates to give  $H^+$  most easily

[MP PET 1994]

- (a)  $H_2O$
- (b)  $H_2S$
- (c)  $H_2Te$
- (d)  $H_2Se$
- **72.** Superphosphate is the mixture of **[CPMT 1993]** 
  - (a) Calcium phosphate and dil.  $H_2SO_4$
  - (b) Sodium phosphate and dil.  $H_2SO_4$
  - (c) Potassium phosphate and dil.  $H_2SO_4$
  - (d) None of these
- **73.** Among  $KO_2$ ,  $NO_2^-$ ,  $BaO_2$  and  $NO_2^+$  unpaired electron is present in [IIT 1997]
  - (a)  $NO_2^+$  and  $BaO_2$
- (b)  $KO_2$  and  $BaO_2$
- (c)  $KO_2$  only
- (d)  $BaO_2$  only
- 74. Which is not easily soluble in water [CPMT 1994]
  - (a)  $H_2$
- (b)  $O_2$
- (c) SO<sub>2</sub>
- (d) CO<sub>2</sub>
- **75.** Point out in which of the following properties oxygen differs from the rest of the members of its family (Group-VIA)

[MP PMT 1997]

- (a) High value of ionisation energies
- (b) Oxidation states (2, 4, 6)
- (c) Polymorphism
- (d) Formation of hydrides

	/00 s and p-bloc	CK LICITICITES			
<b>76.</b>	In the preparation of s	sulphuric acid, $V_2O_5$ is used		(c) Se	(d) Na
		is [CBSE PMT 2001; AFMC 2001]	88.		g is a suboxide [DPMT 2001]
	(a) $S + O_2 \rightarrow SO_2$			(a) $Ba_2O$	(b) <i>Pb</i> <sub>2</sub> <i>O</i>
	(b) $2SO_2 + O_2 \rightarrow 2SO_3$			(c) $C_3O_2$	(d) <i>ZnO</i>
	(c) $SO_2 + H_2O \rightarrow H_2SO_2$	4	89.		f sulphuric acid by contact
	(d) $N_2 + 3H_2 \rightarrow 2NH_3$			process, Tyndall box is	
77•		ng hydrides has the lowest		(a) Filter dust particle	
	boiling point			(b) Remove impurities	
	(a) $H_2O$	[MP PET 1997] (b) $H_2S$		(c) Convert $SO_2$ to $SC_2$	
	<del>-</del>	<del>-</del>		(d) Test the presence	-
-0	(c) $H_2Se$	(d) $H_2Te$	90.		d is known as[Bihar CEE 1995]
78.		the manufacture of $H_2SO_4$ by		(a) Marshell's acid	(b) Caro's acid
	contact process is (a) $Al_2O_3$	[UPSEAT 1999] (b) $Cr_2O_3$	91.	(c) Sulphuric acid $KO_2 + CO_2 \rightarrow ?$ (gas)	(d) None of these [CPMT 1997]
	(c) $V_2O_5$	(d) $MnO_2$	91.		
	- *	<del>-</del>		(a) $H_2$ (c) $O_2$	(b) $N_2$
79.		The molecular formula of sulphur is [MP PMT 1996; MP PET/PMT 1998]			(d) <i>CO</i>
	(a) $S$ (b) $S_2$			$H_2SO_4$ acts as dehyd:	rating agent in its reaction
	(c) $S_4$	(d) $S_8$		with	
80.	•	g is not suitable for use in a			[JIPMER 2001]
00.	descicator to dry subst		ſ.	(a) $H_2C_2O_4$ AIIMS 1996]	(b) $Ba(OH)_2$
	•	(b) $Na_2SO_4$	_	(c) <i>KOH</i>	(d) <i>Zn</i>
	(c) CaCl <sub>2</sub>	(d) $P_4O_{10}$	93.	Which of the followir boiling point	ng group shows the highest
81.	Which shows polymorp	phism [DCE 2000]			[MP PET 2002]
	(a) O	(b) S		(a) $H_2O$	(b) $H_2S$
	(c) Se	(d) All		(c) <i>H</i> <sub>2</sub> <i>Se</i>	(d) $H_2Te$
82.	All the elements of oxy	gen family are [MP PET/PMT 19	981 94.	In presence of moistur	re, <i>SO</i> <sub>2</sub> can <b>[BVP 2003]</b>
	(a) Non-metals	(b) Metalloids		(a) Act as oxidant	(b) Lose electron
00	(c) Radioactive	(d) Polymorphic		(c) Gain electron	(d) Not act as reductant
83.	known as	s of elemental oxygen is	95.	A gas that cannot be co	
	miowii do	[Kerala (Med.) 2002]			[Kurukshetra CEE 1998]
	(a) Azone	(b) Polyzone		(a) $N_2$	(b) O <sub>2</sub>
	(c) Triozone	(d) Ozone		(c) $SO_2$	(d) $PH_3$
84.	When $H_2S$ gas is pas product is	sed through nitric acid, the	96.	Which of the following water on sodium perox	g is formed by the action of
		[Kerala (Engg.) 2002]		(a) $H_2$	(b) $N_2$
	(a) Rhombic S	(b) Prismatic S		(c) $O_2$	(d) $CO_2$
	(c) Amorphous S	(d) Monoclinic S	0=		
	(e) None of these	-	97.	Sulphur on boiling wit	th <i>NaOH</i> solution gives [Roorkee 1999]
85.	Shape of $O_2F_2$ is similar	ar to that of [AIIMS 2004]		(a) $Na_2S_2O_3 + NaHSO_3$	
	(a) $C_2F_2$	(b) $H_2O_2$			(d) $Na_2SO_3 + SO_2$
	(c) $H_2F_2$	(d) $C_2H_2$		2 0 2	2 0 2
86.	Which of the followi	ng bonds has the highest	98.	Quartz is a crystalline	
	energy			(a) Silicon carbide	(b) Sodium silicate
		[CBSE PMT 1996]		(c) Silica	(d) Silicon
	(a) <i>Se</i> − <i>Se</i>	(b) $Te - Te$	99.		gent for the absorption of
_	(c) $S-S$	(d) <i>O</i> – <i>O</i>		$SO_3$ is	Perre 101 elle abbot beton 01
87.	_	g is not a chalcogen[CPMT 1999]		J.	[BHU 2004; DPMT 2004]
	(a) <i>O</i>	(b) S		(a) $80\% H_2SO_4$	(b) $98\% H_2SO_4$
				2 T	<b>≟</b> ₹

(c)  $50\% H_2SO_4$ (d)  $20\% H_2S_2O_7$ **100.** Conc.  $H_2SO_4$  is diluted [Pb. CET 2001] (a) By adding water in  $H_2SO_A$ (b) By adding  $H_2SO_4$  in water (c) By adding glacial acetic acid in  $H_2SO_4$ (d) None of the above 2. **101.** The smog is essentially caused by the presence of [AIEEE 2004] (a) Oxides of sulphur and nitrogen (b)  $O_2$  and  $N_2$ 3. (c)  $O_2$  and  $O_3$ (d)  $O_3$  and  $N_2$ **102.** Bleaching action of  $SO_2$  is due to its [CPMT 2004] (a) Oxidising property (b) Acidic property (d) Reducing property (c) Basic property 103. Oxygen is not evolved on reaction of ozone with [Pb. PMT 2004] (a)  $H_2O_2$ (b) SO, (c) Ha (d) KI **104.** When  $PbO_2$  reacts with conc.  $HNO_3$  the gas 5. evolved is [IIT-JEE (Screening) 2005] (a)  $NO_2$ (b)  $O_2$ (d)  $N_2O$ (c)  $N_2$ 105. Sulphur in +3 oxidation state is present in [DCE 2003] 6. (b) Pyrosulphuric acid (a) Sulphurous acid (d) Thiosulphuric acid (c) Dithionous acid **106.**  $SO_2 + H_2S \rightarrow \text{product.}$  the final product is [Orissa JEE 2005] 7. (a)  $H_2O + S$ (b)  $H_2SO_4$ (d)  $H_2S_2O_3$ (c)  $H_2SO_3$ **107.** Oncontrolled hydrolysis and condensation,  $R_3SiCl$ 8. vields [Orissa JEE 2005] (a)  $R_3Si - O - SiR_3$ (b)  $\{R_3Si - O - SiR_3\}$ 9. R R -Si - O - Si -(c)  $R_3SiOH$ 0 o-Si - O - Si -10. 108. Ozone deplict due to the formation of following compound in Antarctica [Kerala CET 2005] (a) Acrolein (b) Peroxy acetyl nitrate Which one of the halogen acids is a liquid[MP PMT 1985] 11. (c)  $SO_2$  and  $SO_3$ (d) Chlorine nitrate (a) *HF* (e) Formaldehyde (c) HBr

Halogen family

s and p-Block Elements 789 The correct order of the thermal stability of hydrogen halides (H - X) is [AIEEE 2005] (a) HI > HBr > HCl > HF(b) HF > HCl > HBr > HI(c) HCl < HF < HBr < HI(d) HI > HCl < HF < HBrPhosgene is the common name of [CPMT 1974, 86; DPMT 1989; MP PMT 1994] (a) Carbonyl chloride (b) Phosphine (c) Phosphorus oxychloride (d)Phosphorus trichloride The solubility of iodine in water increases in the presence of [CPMT 1973, 74, 78, 86, 89, 91; NCERT 1973; AFMC 1995] (a) Alcohol (b) Chloroform (c) Sodium hydroxide (d) Potassium iodide When thiosulphate ion is oxidised by iodine, which one of the following ion is produced [CPMT 1989; AFMC 1990; CBSE PMT 1996] (a)  $SO_3^{2-}$ (b)  $SO_4^{2-}$ (c)  $S_4 O_6^{2-}$  (Tetrathionate) (d)  $S_2O_6^{2-}$ Bromine is liberated when an aqueous solution of potassium bromide is treated with [CBSE PMT 1989] (a)  $Cl_2$ (b)  $I_2$ (c) Dilute  $H_2SO_4$ (d) SO, Which of the following has greatest reducing power [CPMT 1984, 88, 89, 94] (a) HI (b) HBr (c) HCl (d) HF Chlorine was discovered by [BHU 1988] (a) Davy (b) Priestley (c) Rutherford (d) Sheele Bad conductor of electricity is [MP PET/PMT 1988] (a)  $H_2F_2$ (b) HCl (c) HBr (d) HI Which of the following will not occur[MP PET/PMT 1988] (a)  $Fe + H_2SO_4 \rightarrow FeSO_4 + H_2$ (b)  $Cu + 2AgNO_3 \rightarrow Cu(NO_3)_2 + 2Ag$ (c)  $2KBr + I_2 \rightarrow 2KI + Br_2$ (d)  $CuO + H_2 \rightarrow Cu + H_2O$ Bromine is obtained on commercial scale from [CPMT 1985] (a) Caliche (b) Carnellite (c) Common salt (d) Cryolite

(b) HCl

(b) HBr

(d) HCl

[MP PMT 1985]

(d) HI

Which one of the following acids is the weakest

12.

(a) HClO

(c) HClO<sub>3</sub>

13.	Chlorine reacts with various condition to give	sodium hydroxide under ve	25.[]	(d) Bromine is more re NCWNE119731r is treated		
	(a) Sodium chloride	(b) Sodium hypochlorite		redish brown gas evol		[Pb. CET 2003]
	(c) Sodium chlorate	(d) All of these		(a) Mixture of broming	_	[FU. CET 2003]
14.	$Br_2$ gas turns starch io			(b) HBr	z anu mbi	
	-	1987; AFMC 1987; AMU 1999]		(c) Bromine		
	(a) Blue	(b) Red		(d) None of these		
	(c) Colourless	(d) Yellow	26.	Sea weed is employed	ac a cource	of manufacture
15.		sed through hot and conc.	20.	of	as a source	or manufacture
٠,٠		wing compound is formed			1982, 86, 200	2; MP PET 2002]
	solution of Koll, lono	[CPMT 1971, 79; BVP 2003]		(a) <i>F</i>	(b) <i>I</i>	,
	(a) KCl	(b) <i>KClO</i> <sub>3</sub>		(c) Br	(d) <i>Cl</i>	
		-	27.	Which of the following		ive halogen
	(c) KClO <sub>2</sub>	(d) KClO <sub>4</sub>	,	•	,	[MP PET 1990]
16.	Deacon's process is use	ed in the manufacture of		(a) $Cl_2$	(b) $Br_2$	
	(a) Blacking novedon	[BHU 1979]		(c) $I_2$	(d) $F_2$	
	(a) Bleaching powder	(b) Sulphuric acid	20	<del>-</del>	-	mmocomto alcon
10	<ul><li>(c) Nitric acid</li><li>Which of the following</li></ul>	(d) Chlorine	28.	Which of the for electropositive properti	•	presents clear
17.		IT 1988, 2000; MP PMT 1995;				8; MP PMT 1991]
		), 90; Kurukshetra CEE 1998;		(a) <i>F</i>	(b) <i>Cl</i>	-, <b>55</b> - <u>-</u>
		Roorkee 2000; RPMT 2000]		(c) Br	(d) <i>I</i>	
	(a) HF	(b) HCl	29.	Which statement is fa	lse	[MP PET 1991]
	(c) HBr	(d) <i>HI</i>		(a) Electronegativity	of fluorine is	maximum
18.	Which is the most vola			(b) Electron affinity of	f fluorine is 1	maximum
		979; AIIMS 1980; DPMT 1982;		(c) Melting point of flu		
	(a) HF	J & K CET 2005; DPMT 2002] (b) <i>HCl</i>		(d) Boiling point of flu		
	(c) HBr	(d) <i>HI</i>	30.	Strongest reducing ag		[MP PMT 1990]
19.		s solution of KClO <sub>3</sub> with		(a) $F^{-}$	(b) <i>Cl</i> <sup>-</sup>	
_5:	iodine the following n	roduct is obtained[NCFRT 1080	<b>5</b> 1	(c) Br <sup>-</sup>	(d) $I^-$	
	iodine, the following product is obtained[NCERT 198 (a) KIO <sub>3</sub> (b) KCIO <sub>4</sub>			Which of the follow		
	(c) $KIO_4$	(d) KCl		shell electronic config		-
	·			[MP PET 1991; Man		5; MP PM1 1996]
20.	Colour of iodine solution is disappeared by shaking it with aqueous solution of [CPMT 1979, 81; N			(a) $s^2 p^3$	(b) $s^2 p^6$	
	MP	9 301011011 01[CFM1 1979, 81, M PMT 1986; RPMT 1997, 2002]	F FEI,	(c) s p	(d) $s^2 p^5$	
	(a) $H_2SO_A$	(b) $Na_2S$	32.	Chlorine can remove		[MP PET 1990]
	(c) $Na_2S_2O_3$	(d) $Na_2SO_4$		(a) Br from NaBr sol		
<b>0</b> 4		= :		(b) F from NaF solution		
21.		$l_2$ gas may be made by		(c) Cl from NaCl solu		
		MnO <sub>4</sub> with a concentrated		(d) $F$ from $CaF_2$ solut		
	solution of	[CDMT 10=0]	33.	Hydrolysis of which occur	of the folio	owing does not
	(a) Potaccium chlorido	[CPMT 1973] (b) Sodium chloride		occur		[AIIMS 1982]
	<ul><li>(a) Potassium chloride</li><li>(c) Bleaching powder</li></ul>	(d) Hydrochloric acid		(a) VCl <sub>4</sub>	(b) TiCl <sub>4</sub>	[
22.		ongst the following is[IIT 1989]	1	(c) SiCl <sub>4</sub>	(d) <i>CCl</i> <sub>4</sub>	
22.	(a) HClO <sub>4</sub>	(b) $HClO_3$		Nitric acid converts io	·	[MD DMT 1000]
	·	-	34.	(a) Iodic acid		[MP PMT 1990] piodic acid
	(c) $HClO_2$	(d) HClO		(c) Iodine nitrate	-	e pentaoxide
23.	Iodine deficiency in die		35.	In $KI$ solution, $I_2$ rea		
	(a) Nightblindness	(b) Rickets				9; EAMCET 1992]
	(c) Goitre	(d) Beri-beri		(a) $I^-$	(b) $KI_2$	, EAMCEI 1992]
24.	Which of the following	is correct [CPMT 1985]			=	
	(a) Iodine is a solid	o in victor		(c) $KI_2^-$	(d) $KI_3$	
	(b) Chlorine is insolubl		36.	Which reaction canno	t be used for	=
	(c) Iodine is more reac	tive than bromme		of halogen acid		[MP PMT 1989]

			s and p-Block Elements 791
	(a) $2KBr + H_2SO_4 \rightarrow K_2SO_4 + 2HBr$		(c) Br (d) I
	(b) $NaHSO_4 + NaCl \rightarrow Na_2SO_4 + HCl$	46.	Mark the element which shows only one
	(c) $NaCl + H_2SO_4 \rightarrow NaHSO_4 + HCl$		oxidation state
			[BHU 1988; MP PET 2002]
	(d) $CaF_2 + H_2SO_4 \rightarrow CaSO_4 + 2HF$		(a) $F$ (b) $Cl$
37.	In dark, which of the following reacts with hydrogen		(c) $Br$ (d) $I$
	[MP PMT/PET 1988; Kurukshetra CEE 1998]	47•	Which of the following arrangement for the three
	(a) $Br_2$ (b) $F_2$		halogens <i>Cl</i> , <i>Br</i> and <i>I</i> when placed in the order
	(c) $I_2$ (d) $Cl_2$		of their increasing electron affinity is correct[CPM
38.	The more activeness of fluorine is due to[MP PMT 199]	001	(a) Cl, Br, I (b) I, Br, Cl
50.	(a) <i>F-F</i> bond has less energy		(c) Br, Cl, I (d) I, Cl, Br
	(b) $F_2$ is gas at normal temperature	48.	8 8
	(c) Its electronic bond is maximum		agent
	(d) <i>F-F</i> bond has more energy		[CPMT 1978, 91, 94; MNR 1990; AMU 1983, 84;
39.	Which of the following after reacting with <i>KI</i> do		MP PMT 1991, 92, 96; IIT 1992; UPSEAT 2000]
	not remove iodine [MP PET 1989]		(a) $F_2$ (b) $Cl_2$
	(a) $CuSO_4$ (b) $K_2Cr_2O_7$		(c) $Br_2$ (d) $I_2$
	(c) $HNO_3$ (d) $HCl$	49.	Fluorine is a better oxidising agent than $Br_2$ . It is
40.	Aqueous solution of which of the following acids		due to
_	cannot be kept in a bottle of glass [MP PET 1989]		[EAMCET 1992]
	(a) HF (b) HCl		(a) Small size of fluorine
	(c) HBr (d) HI		(b) More electron repulsion in fluorine
41.	Which of the following pairs is not correctly		(c) More electronegativity of fluorine
	matched		(d) Non-metallic nature of fluorine
	[MP PET 1993]  (a) A halogen which is liquid at room temperature—Bromine	50.	Fluorine is a stronger oxidising agent than chlorine in aqueous solution. This is attributed to many factors except
	(b) The most electronegative element— <i>Fluorine</i>		[Pb. CET 1989]
	(c) The most reactive halogen—Fluorine		(a) Heat of dissociation (b) Electron affinity
	(d) The strongest oxidizing halogen— <i>Iodine</i>		(c) Heat of hydration (d) Ionisation potential
42.	Iodine is formed when potassium iodide reacts with a solution of [MNR 1984; MP PET/PMT 1998]	51.	Mark the element which displaces three halogens from their compounds
	(a) $ZnSO_4$ (b) $CuSO_4$		[MP PMT 1980, 82; BHU 1984; NCERT 1987]
	(c) $(NH_4)_2SO_4$ (d) $Na_2SO_4$		(a) $F$ (b) $Cl$
42	As the atomic number of halogens increases, the		(c) Br (d) I
43.	halogens	52.	Which one of the following is the most basic
	[MP PMT 1991]		[CPMT 1975, 77; MP PMT 2001] (a) I (b) Br
	(a) Lose the outermost electrons less readily		(a) <i>I</i> (b) <i>Br</i> (c) <i>Cl</i> (d) <i>F</i>
	(b) Become lighter in colour	53.	Which of the following will displace the halogen
	(c) Become less denser	55.	from the solution of the halide [EAMCET 1979]
	(d) Gain electrons less readily		(a) $Br_2$ added to $NaCl$ solution
44.	Which statement is correct about halogens [EAMCET 1991]		(b) $Cl_2$ added to $KCl$ solution
	(a) They are all diatomic and form univalent ions		(c) KCl added to NaF solution
	(b) They are all capable of exhibiting several oxidation states	54.	(d) $Br_2$ added to $KI$ solution Fluorine does not form positive oxidation states
	(c) They are all diatomic and form divalent ions		because
	(1) The control of the state of		[AIIMS 1987]

(a) It is most electronegative element(b) It forms only anions in ionic compounds

(c) It cannot form multiple bonding

(d) They can mutually displace each other from the solution of their compounds with metals

(b) *Cl* 

[CPMT 1984, 89]

Mark the smallest atom

45.

(a) F

(d) It shows non-bonded electron pair repulsion (a) Killing germs (b) Prevention Ωf pollution due to small size In the isolation of fluorine, a number of (c) Cleansing (d) Removing dirt 55. difficulties were encountered. Which statement is Chlorine cannot be used [MP PET/PMT 1988] 65. correct (a) As bleaching agent [NCERT 1983, 86] (b) In sterilisation (a) The potential required for the discharge of the (c) In preparation of antiseptic fluoride ions is the lowest (d) For extraction of silver and copper (b) Fluorine reacts with most glass vessels Chlorine acts as a bleaching agent only in 66. (c) Fluorine has great affinity for hydrogen presence of (d) Electrolysis of aqueous HF gives ozonised [IIT 1983; DCE 2002] oxygen (a) Dry air (b) Moisture Fluorine reacts with water to give [BHU 1988, 89] 56. (c) Sunlight (d) Pure oxygen (a) HF and  $O_2$ (b) HF and  $OF_2$ 67. Euchlorine is a mixture of [CPMT 1988] (a)  $Cl_2$  and  $SO_2$ (b)  $Cl_2$  and  $ClO_2$ (c) HF and  $O_3$ (d)  $HF, O_2$  and  $O_3$ (c)  $Cl_2$  and CO(d) None of these In which of the following, oxygen has + 2 57. oxidation number [EAMCET 1986] A gas reacts with CaO, but not with NaHCO<sub>3</sub>. 68. (a)  $F_2O$ (b) Cl<sub>2</sub>O The gas is [AFMC 1987] (c)  $Na_2O_2$ (d)  $Na_2O$ (a) CO<sub>2</sub> (b) Cl<sub>2</sub> The electrolysis of a certain liquid resulted in the 58. (c)  $N_2$ (d)  $O_2$ formation of hydrogen at the cathode and chlorine at the anode. The liquid is When FAMOFTA 939 assed over dry slaked lime at 69. (a) Pure water room temperature, the main reaction product is (b)  $H_2SO_4$  solution [CBSE PMT 1992] (a)  $Ca(ClO_2)_2$ (b) *CaCl*<sub>2</sub> (c) NaCl solution in water (d) CuCl<sub>2</sub> solution in water (c) CaOCl<sub>2</sub> (d)  $Ca(OCl_2)_2$ In the preparation of chlorine from HCl, MnO<sub>2</sub> 70. Bromine is obtained commercially from sea water 59. by adding [CPMT 1988] acts as (a)  $AgNO_3$  solution (b) Crystals of *NaBr* [CPMT 1981] (b) Reducing agent (c) Cl, (a) Oxidising agent (d)  $C_2H_4$ (c) Catalytic agent (d) Dehydrating agent In the manufacture of bromine from sea water, 71. Chlorine gas is dried over 60. [CPMT 1980] the mother liquor containing bromides is treated (a) *CaO* (b) NaOH [CBSE PMT 1992; MP PMT 2001; BHU 2002; (c) KOH (d) Conc.  $H_2SO_4$ JIPMER 2002] Chlorine can be manufactured from [CPMT 1989] 61. (a) CO<sub>2</sub> (b) Cl<sub>2</sub> (a) Electrolysis of NaCl (b) Electrolysis of brine (c)  $I_2$ (d) SO<sub>2</sub> (c) Electrolysis of bleaching powder  $Br^-$  is converted into  $Br_2$  by using 72. [CPMT 1987] (d) All of these (a)  $Cl_2$ (b) Conc. HCl 62. When chlorine water is exposed to sunlight,  $O_2$  is (c) HBr (d)  $H_2S$ [AFMC 1989] liberated. Hence (a) Hydrogen has little affinity to  $O_2$ A salt, which on heating with conc.  $H_2SO_4$  gives (b) Hydrogen has more affinity to  $O_2$ violet vapours, is [CPMT 1971] (a) Iodide (b) Nitrate (c) Hydrogen has more affinity to  $Cl_2$ (c) Sulphate (d) Bromide (d) It is a reducing agent When  $I_2$  is dissolved in  $CCl_4$ , the colour that When cold NaOH reacts with Cl2 which of the 63. results is following is formed [AFMC 1992] [AFMC 1993] (a) NaClO (b) NaClO, (a) Brown (b) Violet (c) NaClO<sub>3</sub> (d) None of these (c) Colourless (d) Bluish green Chlorine is used in water for [CBSE PMT 1988] 64. Which of the following halogen oxides is ionic[CPMT 1989

				s and p-B	Block Elements 793
	(a) ClO <sub>2</sub>	(b) <i>BrO</i> <sub>2</sub>		•	NCERT 1974; CPMT 1976, 90]
	(c) $I_2O_5$	(d) $I_4 O_9$		(a) <i>HF</i>	(b) <i>HCl</i>
G	- *	. ,		(c) HBr	(d) <i>HI</i>
6.	KI when heated with o	- '	86.		wing hydrogen halides has the
	(2) 111	[MP PET/PMT 1988]			nt[AIIMS 1980; DPMT 1983; MP PM
	(a) <i>HI</i>	(b) I <sub>2</sub>		(a) <i>HF</i> (c) <i>HBr</i>	(b) <i>HCl</i> (d) <i>Hi</i>
	(c) $HIO_3$	(d) $KIO_3$	87.	Hydrogen bonding i	
7•	• • • • • • • • • • • • • • • • • • • •	HCl molecule is[AIIMS 1992]	07.		1989; DPMT 1990; Roorkee 1995]
	(a) Pure covalent	(b) Polar covalent		(a) <i>HF</i>	(b) HCl
_	(c) Highly covalent	(d) <i>H</i> -bonding		(c) HBr	(d) <i>HI</i>
8.	is because	is a low boiling liquid. This [EAMCET 1981, 89]	88.	The type of hybrid in $ClO_2^-$ is	orbitals used by chlorine atom
	(a) $H - F$ bond is stron			(a) $SP^{3}$	$(h)$ $SD^2$
	(b) $H - F$ bond is weak			(a) <i>SP</i> (c) <i>SP</i>	(b) SP <sup>2</sup> (d) None of these
		regate because of hydrogen	89.	Which one is the an	
	bonding in HF		og.		IIINS 1983; BHU 1983; AMU 1984]
	(d) HF is a weak acid	ad by the action of the		(a) $Cl_2O$	(b) ClO <sub>2</sub>
9.	$H_2SO_4$ on $KI$ because	ed by the action of conc. [MNR 1984]		(c) $Cl_2O_6$	(d) $Cl_2O_7$
			00		ving halogens is solid at room
	(a) <i>HI</i> is stronger than		90.	temperature	[MP PET 1999; AFMC 1999]
	(b) <i>HI</i> is more volatile	than $H_2SO_4$		(a) Chlorine	(b) Iodine
	(c) $H_2SO_4$ is an oxidist	ing agent		(c) Bromine	(d) Fluorine
	(d) $H_2SO_4$ forms comp	olex	91.	Which of the follow	ring chemical contains chlorine
).		water is good conductor			[MP PET 1999]
-		en chloride is not. This is		(a) Fischer salt	(b) Epsom salt
	due to the reason that	[NCERT 1980]		(c) Fremy's salt	(d) Spirit of salt
	(a) Water is a good con	ductor of electricity	92.	The element which in a chemical reacti	never acts as reducing agent on is [Bihar CEE 1995]
	(b) HCl in water ionise	es		(a) O	(b) <i>Li</i>
	(c) Gas can not conduc	t electricity but water can		(a) <i>U</i> (c) <i>F</i>	(d) C
	(d) None of these		93.		reacts with $I_2$ to give
•	Sodium chloride when	heated with conc. $H_2SO_4$		-	89; Roorkee 1995; JIPMER 2001]
	-	chromate gives[CPMT 1981, 84]	]	(a) <i>HI</i>	(b) <i>HOI</i>
	(a) Chromic chloride	(b) Chromyl chloride		(c) <i>HOIO</i> <sub>3</sub>	(d) HOIO <sub>3</sub>
2.		(d) None of these oes not play any role in	94.	The formula of sor	me fluorides are given below. combine further with fluorine[N
	boiling of			(a) $IF_5$	(b) <i>NaF</i>
	(a) MI	[AFMC 1992]		(c) <i>CaF</i> <sub>2</sub>	(d) $SF_5$
	(a) $NH_3$	(b) $H_2O$	95.	Which one below is	5
	(c) HI	(d) $C_2H_5OH$	93•	(a) $CN^-$	(b) <i>ICl</i>
3.		sulphuric acid, HCl can			
	reduce $KMnO_4$ and $HF$	can reduce[IIT 1981; MP PET	_	(c) <i>IF</i> <sub>5</sub>	(d) $I_3^-$
	(a) $H_2SO_4$	(b) $KMnO_4$	96.	Which one is highes	
	(c) $K_2Cr_2O_7$	(d) None of these		(a) $N_{\alpha}C^{\dagger}$	[AIIMS 1982; BHU 1999]
1.	Which has the highest	molar heat of vaporisation		(a) NaCl	(b) <i>NaBr</i>
	-	[CPMT 1991]	07	(c) NaF	(d) NaI
	(a) <i>HF</i>	(b) HCl	97.	halide has	is correct because the chosen
	( ) IID	( 1) ****			

[AIIMS 1982]

(a) Minimum ionic character

(b) Maximum ionic character

(c) HBr

**85.** Mark the strongest acid

(d) HI

[Bihar MEE 1996; MP PET/PMT 1998;

#### 794 s and p-Block Elements (c) Highest oxidising power 109. Which of the following statements is not true [CBSE PMT 2003] (d) Lowest polarity (a) HF is a stronger acid than HCl Which of the following oxidizes $H_2O$ to oxygen 98. (b) Among halide ions, iodide is the most [MP PET 1994] powerful reducing agent (a) Chlorine (b) Fluorine (c) Fluorine is the only halogen that does not (c) Bromine (d) Iodine show a variable oxidation state The bleaching action of the bleaching powder is 99. (d) HOCl is a stronger acid than HOBr due to the liberation of [MP PMT 1994] 110. The correct order of acidic strength [Pb. CET 2004] (a) Chlorine (b) Molecular oxygen (a) $Cl_2O_7 > SO_2 > P_4O_{10}$ (b) $K_2 > CaO > MgO$ (c) Nascent oxygen (d) Calcium carbonate (c) $CO_2 > N_2O_5 > SO_3$ (d) $Na_2O > MgO > Al_2O_3$ 100. Which of the following element is extracted 111. Halogen acid used in the preparation of aqua commercially by the electrolysis of an aqueous regia is solution of its compound [DPMT 2002] [KCET 2002] (a) HBr (b) HI (b) Bromine (a) Chlorine (c) HCl (d) HF (d) Calcium (c) Aluminium 112. NaOCl is used as a bleaching agent and sterilising **101.** The effective component of bleaching powder is ..... agent. It can be synthesized by the action of [RPET 2003] of calcium (a) NaCl with $H_2O$ [Kerala (Engg.) 2002] (b) $NH_4Cl$ with NaOH(a) Chlorine (b) Bromine (c) Aluminium (d) Calcium (c) $Cl_2$ with cold and dilute NaOH 102. $Na_2S_2O_3 + I_2 \rightarrow \text{Product is}$ [BHU 2003] (d) Cl2 with hot and concentrated NaOH (a) $Na_2S$ (b) NaI 113. Metal halide which is insoluble in water is[AIIMS 1996] (c) $Na_2S_4O_6$ (d) $S_2$ (a) AgI (b) *KBr* 103. Which of the following is prepared by electrolytic (c) *CaCl* , (d) AgF method 114. Which one among the following non-metals liquid [CBSE PMT 2001] at 25°C (a) Ca (b) Sn [MP PMT 1999] (c) S (d) $F_2$ (a) Bromine (b) Carbon 104. Beilstein test is used for [AFMC 1995] (c) Phosphorus (d) Sulphur (a) $N_2$ (b) Cl 115. Bleaching action of chlorine is due to [Bihar CEE 1995] (a) Oxidation (b) Reduction (c) Na (d) CO<sub>2</sub> (c) Hydrolysis (d) Its acidic nature **105.** Which one will liberate $Br_2$ from KBr[BVP 2003] 116. Hydrogen iodide cannot be prepared by the action (b) SO<sub>2</sub> (a) $I_2$ of conc. $H_2SO_4$ on potassium iodide because [Bihar CEE 19] (c) HI (d) Cl<sub>2</sub> (a) HI is stronger than $H_2SO_4$ 106. Chlorine dioxide is best prepared by passing dry (b) HI is more volatile than $H_2SO_4$ [Kerala PMT 2003] (a) Chlorine gas over hot HgO(c) $H_2SO_4$ is an oxidising agent (b) Chlorine and oxygen gas over hot pt catalyst (d) $H_2SO_4$ forms complex (c) Hydrogen chloride and oxygen over silver oxide 117. White enamel of our teeth is [Bihar CEE 1995] (d) Hydrogen chloride over phosphorus pentoxide (b) *CaF*<sub>2</sub> (a) $Ca_3(PO_4)_2$ (e) Chlorine over hot silver chlorate (c) *CaCl*<sub>2</sub> (d) $CaBr_2$ 107. The mixture of concentrated HCl and $HNO_3$ made 118. The least active halogen with hydrogen is[DPMT 1996] in 3: 1 ratio contains [AIIMS 2003] (a) Cl (b) I (b) NOCl (a) ClO<sub>2</sub> (c) Br (d) F (c) $NCl_3$ (d) $N_2O_4$ 119. Iodine dissolves readily in [BHU 1996] 108. On exciting $Cl_2$ molecule by UV light, we get (a) Water (b) Potassium iodide [UPSEAT 2003] (c) Carbon tetrachloride (d) Alcohol

(b)  $Cl^+$ 

(d) All

(a) *Cl* (c) *Cl* 

120.	Which one of the follow solution gives a white				(b) All halogens form of (c) All halogens except	=	oxy acids
	acid				(d) Only iodine form oxy acids		
			AMCET 1997]	131.	When iodine reacts with	-	and <i>NaCl</i>
	(a) NaCl	(b) <i>KCl</i>		1310	VVIIOII IOUIIIO I CUCES VVII	i i i i i i i i i i i i i i i i i i i	[CPMT 1997]
	(c) $MgCl_2$	(d) $FeCl_3$			(a) It gives mixture of	$F_{s}$ $Cl_{s}$ and $R$	
21.	Which of the following					2, 612 0110 2	• 2
	reference to the oxidati			1997]	(b) It gives chlorine		
	(a) $I_2 > ICl < HI < HIO_4$	(b) $HIO_4 < IC$	$l < I_2 < HI$		<ul><li>(c) It gives bromine</li><li>(d) None of these</li></ul>		
	(c) $I_2 < HI < ICl < HIO_4$	(d) $HI < I_2 < I_2$	$ICl < HIO_4$	122	Which is the strongest of	of the followin	ng acidell <b>ipme</b> p
22.	The correct order of ac	dic strength is		132.	(a) $HClO_4$	(b) $H_2SO_4$	ig acidstiffmen
[P	b. PMT 1998; AFMC 1998;1	KCET 2000; Oris	sa JEE 2005]		·		
	(a) $HF < HCl < HBr < HI$				(c) HCl	(d) $HNO_3$	
	(b) $HCl < HBr < HF < HI$		133.	Hydrogen has a tenden			
	(c) $HBr < HCl < HI < HF$				acquire helium configuration. In this respect it resembles		
	(d) $HI < HBr < HCl < HF$				resembles		[JIPMER 1999]
23.	The chief source of iodine in which it is present				(a) Halogens	(b) Actinide	
	as sodium iodate is	.5 557.2			(c) Transition elements		
	(a) Sea weeds			134.	What is the product of		
	(b) Caliche				$HgCl_2$ and $Hg(CN)_2$		MP PET 2002]
	(c) Carnallite				(a) $(CN)_2$		
	(d) Iodine never exists				(b) Addition compound	HaCl Ha(CN)	
24.	The lattice energy of the following order	ne lithium halio [Roorkee Qua]			(c) $Hg(CN)Cl$	118012.118(014)	2
	(a) $LiF > LiCl > LiBr > Li$	!			(d) $Hg[Hg(CN)_2Cl_2]$		
	(b) $LiCl > LiF > LiBr > LiBr$	!		135.	The weakest acid HX (X	f = F, Cl, Br, I	is[BHU 2000]
	(c) $LiBr > LiCl > LiF > Li$	!		-33.	(a) <i>HF</i>	(b) <i>HCl</i>	10[2110 2000]
	(d) $LiI > LiBr > LiCl > LiF$	7			(c) HBr	(d) <i>HI</i>	
25.	Which has the strong bo	ond	[DCE 2001]	136.	Bleaching powder is ob	tained by pas	ssing chlorine
	(a) <i>F</i> – <i>F</i>	(b) <i>F - Cl</i>			on		
	(c) $F - Br$	(d) <i>Cl</i> – <i>B</i>					[KCET 2002]
<b>26</b> .	Iodine and hypo react to	produce	[DPMT 2001]		(a) Lime stone	(b) Quick li	
	(a) $Na_2S$	(b) $Na_2SO_4$			(c) Slaked lime	(d) Pure lin	
	(c) $Na_2S_4O_6$	(d) $Na_2SO_3$		137.	Chlorine is liberated, w		
27	Bleaching powder is con		nted as[RDMT 14	1700	(a) $KMnO_4 + NaCl$	(b) $K_2 C r_2 O_7$	$+MnO_2$
-/•	(a) CaOCl <sub>2</sub>	(b) CaO	.cou as[KFWI I	1/50	(c) $Pb_2(NO_3)_4 + MnO_2$	(d) $K_2Cr_2O_7$	+HCl
	(a) CaO(Cl)	(d) CaCl(OCl)		138.	Which of the followin maximum use in photog	-	pounds finds
28.	When chlorine reacts	s with cold	and dilute		(a) AgCl	(b) AgBr	
	solution of sodium						
	obtained are	[CBSE PMT 199	98]		(c) AgI	(d) $AgNO_3$	
	(a) $Cl^- + ClO^-$	(b) $Cl^{-} + ClO_{2}^{-}$		139.	Which of the following positive oxidation state		
	(c) $Cl^- + ClO_3^-$	(d) $Cl^- + ClO_4^-$				[EAMCET 1997	7; AIIMS 2000]
29.	A one litre flask is full	of brown brom	ine vapour.		(a) <i>Cl</i>	(b) <i>Br</i>	
	The intensity of brown	colour of vapo	our will not		(c) I	(d) F	
	decrease appreciably or			E14467	<b>1998</b> Istrength of oxy ac	ids of chlorin	e follows the
	(a) Pieces of marble	(b) Carbon di	sulphide		order		
	(c) Carbon tetrachlorid	e (d) Animal	charcoal				SSE PMT 2005]
owo					(a) $HClO < HClO_2 < HClO_3$	$O_3 < HClO_4$	
30.	Which of the following  (a) Only chlorine and by			7]	(b) $HClO_4 < HClO_3 < HC$	$lO_2 < HClO$	

	796 s and p-Block	k Elements					
	(c) $HClO_4 < HClO_3 < HClO_3 < HClO_4$	ClO < HClO <sub>2</sub>			(c) CaBr <sub>2</sub>	(d) <i>CaI</i> <sub>2</sub>	
	(d) None of these			151.	Which one of the f		
141.	Bleaching powder is obwith	otained by tro	eating chlorine		when mixed, prod temperature	luces chlorine [IIT 1995]	_
			[Pb. PMT 1999]		(a) $NaCl$ and $MnO_2$		
	(a) CaO	(b) <i>CaCO</i> <sub>3</sub>			(b) $NaCl$ and $HNO_3$	(conc.)	
	(c) <i>CaSO</i> <sub>4</sub>	(d) <i>Ca(OH)</i>	)2		(c) $NaCl$ and $H_2SO$	4 (conc.)	
142.	Which statement is not	true	[MP PET 2000]		(d) HCl (conc.) and	KMnO <sub>4</sub>	
	(a) $Ni(CO)_4$ is diamagn	etic		152.	Concentrated $H_2SO$	•	used to prepare
	(b) $BI_3$ is a stronger Le	ewis acid tha	$n BF_3$		HBr from NaBr, bed		1 1
	(c) Graphite conducts e	lectricity wh	ereas diamond		(a) Reduces <i>HBr</i>	(b) Oxidi	ses <i>HBr</i>
	does not	whereas PC	l is inart		(c) Disproportionate	es <i>HBr</i> (d)Re	acts slowly with
142	(d) <i>CCl</i> <sub>4</sub> is hydrolysed			153.	Which of the follow	_	
143.	Bleaching powder loses long time because	s its power of	[KCET 2000]		has doubtful existen (a) $CI_4$	(b) <i>GeI</i> <sub>4</sub>	[IIT 1996]
	(a) It changes into calc	ium hypochl			(c) $SnI_4$	(d) $PbI_4$	
	(b) It changes into cal	cium chloric	le and calcium	154	Chlorine cannot disp	•	[MP PET 1996]
	hydroxide (c) It absorbs moisture			154.	(a) Fluorine from <i>N</i>		
	(d) It changes into cal		le and calcium		(c) Bromine from N		
chlor	_			155.	When fluoride is h	eated with co	onc. $H_2SO_4$ and
144.	The compound which	forms a dat	rive bond with		$MnO_2$ the gas evolve	ed is	[DPMT 2000]
	ammonia		[JIPMER 2001]		(a) $F_2$	(b) <i>SF</i>	
	(a) <i>CCl</i> <sub>4</sub>	(b) <i>BCl</i> <sub>3</sub>	UII MER 2001)		(c) HF	(d) None	
	(c) $MgCl_2$	(d) NaCl		156.	$Cl_2$ reacts with $CS_2$	in presence	of $I_2$ catalyst to
145.	The bleaching action o		powder is due		form		[AFMC 1995]
10	to the formation of	8	[Roorkee 1999]		(a) CHCl <sub>3</sub>	(b) <i>CCl</i> <sub>4</sub>	[AFMC 1995]
	(a) $CaCl_2$	(b) $CaSO_4$			(c) $C_2H_5Cl$	(d) $C_2H_6$	
	(c) HClO	(d) Ca(ClO	3)2	157.	Amongst <i>LiCl</i> , <i>RbCl</i> , <i>E</i>		
146.	Fluorine with dilute Na	_	[MH CET 2000]	-5/-	minimum ionic cha		
	(a) $OF_2$	(b) $O_3$			compounds		_
	(c) $O_2$	(d) HF an	$d O_2$			(I) PLCI	[RPMT 1999]
147.	Which is not oxidised b	y $MnO_2$	[DCE 2003]		(a) LiCl, MgCl <sub>2</sub>	(b) <i>RbCl</i> ,	-
	(a) <i>F</i>	(b) <i>Cl</i>			(c) RbCl,MgCl <sub>2</sub>	(d) MgCl	-
_	(c) I <sub>2</sub>	(d) <i>I</i>	_	158.	Which is formed wh concentrated sodium		act with hot and
148.	Bromine water reacts v				(a) $O_2$	(b) $O_3$	
		(b) $H_2SO_4$			(c) NaO	(d) <i>HF</i>	
		(d) <i>S</i> and	-	159.	Which of the follow	` '	is used to find
149.	Which of the following				atomic $Cl_2$ from mo	olecular ${\it Cl}_2$	[CPMT 1996]
	(a) $2VI + D_m \rightarrow 2VD_m + I$	-	PMT PMT 2002]		(a) High temperatur		
	(a) $2KI + Br_2 \rightarrow 2KBr + I_2$				(b) Low temperature		
	(b) $2H_2O + 2F_2 \to 2HF +$	_			<ul><li>(c) High temperature</li><li>(d) Low temperature</li></ul>	_	
	(c) $2KBr + I_2 \rightarrow 2KI + Br_2$	_		160.	Which one is least b	=	[JIPMER 2000]
	(d) $2KBr + Cl_2 \rightarrow 2KCl + R$	-			(a) $BI_3$	(b) <i>BBr</i> <sub>3</sub>	
150.	Which of the following	has the lowe			(c) BCl <sub>3</sub>	(d) $BF_3$	
	(2) CaF	(b) CaCl	[Roorkee 2000]	161.	On heating $NaCl + R$	9	$H_2SO_4$ , the gas
	(a) $CaF_2$	(b) <i>CaCl</i> <sub>2</sub>					

comes out is

[JIPMER 2000]

				s and n	-Block	Elements 7	707
	(a) O <sub>2</sub>	(b) Cl <sub>2</sub>	171.	Which of the following			
	(c) $CrOCl_2$	(d) $CrO_2Cl_2$	-,	not possible	8	11101000110 10	one or recours,
160	<del>-</del>						[BHU 2002]
102.	Aqua regia is a mixture (a) $3HCl + 1HNO_3$	of [KCET (Med.) 2001] (b) $H_3PO_4 + H_2SO_4$		(a) $OF_4$		(b) $OF_2$	
	_			(c) SF <sub>4</sub>		(d) $O_2F_2$	
163.	Unlike other halogens	(d) $HCl + CH_3COOH$ s fluorine does not show	172.	Iodine is released with	d when	potassium	iodide reacts
	higher oxidation states					[	[UPSEAT 1999]
	(a) It is highly electron	egative		(a) $ZnSO_4$		(b) $CuSO_4$	
	(b) It has no <i>d</i> -orbitals	work amali		(c) FeSO <sub>4</sub>		(d) $(NH_4)_2 Se$	$O_4$
		ble and isoelectronic with	173.	Which of the follo	owing is	s used in the	e preparation
neon		at aleans are sight a saidation				[CI	BSE PMT 1999]
164.	state	ot show variable oxidation		(a) $Only MnO_2$			
	State	[UPSEAT 2003]		(b) $OnlyKMnO_4$			
	(a) $F_2$	(b) <i>Cl</i> <sub>2</sub>		(c) Both $MnO_2$ an	d <i>KMnC</i>	$O_4$	
	(c) $Br_2$	(d) $I_2$		(d) Either $MnO_2$	or <i>KMnO</i>	$O_4$	
165.	<del>-</del>	fumes of $HF$ are removed	174.	Among $Cl^-$ , $Br^-$ , oxidise to dihalog	$I^-$ , the	·	der for being [CPMT 1999]
		[MH CET 2002]		_		(b) $Cl^{-} > Bm$	
	(a) Solid <i>NaF</i>	(b) $H_2$ gas		(a) $I^- > Cl^- > Br^-$			
	(c) Solid KHF <sub>2</sub>	(d) None of these	175	(c) $I^- > Br^- > Cl^-$ On heating $KClO_3$			
166.	Fluorine is prepared by		1/5.				[CPMT 1999]
	(a) Oxidation of <i>HF</i>			(a) $Cl_2O$		(b) ClO <sub>2</sub>	
	(b) Electrolysis of KF			(c) <i>ClO</i> <sub>3</sub>		(d) $Cl_2O_7$	
	(c) Electrolysis of fuse	$d_{KHF_2}$	176.	For which one			properties of holds good[ <b>MP PE</b> T
	(d) Decomposition of H	$IgF_2$		(a) Electron affin		(b) Electron	•
167.	Amongst halogens flu	orine is most oxidising		(c) Atomic radius	-	(d) Boiling	•
	because		177.	Which of the fol			-
	(a) Fluorine has highes		,,	going down from			
	(b) Fluorine is most ele	y for fluorine molecule is		periodic table ?			
lowe		y for fluorine molecule is		( ) 71 ( )		_	MP PMT 1997]
	(d) All are correct			(a) Electronegative	-	(b) Volatile	
168.		es are soluble in water but e	178.	(c) Ionic radius  Among the haloge		(d) Oxidisin	
	(a) It is amphoteric			nitric acid is			[KCET 2004]
	(b) The $Li-F$ bond is h	nighly ionic		(a) Fluorine		(b) Iodine	[KCE1 2004]
	(c) Its lattice energy is	high		(c) Chlorine		(d) Bromine	<u> </u>
	(d) $Li^+$ ion is least hyd		179.	The reaction of th		` '	
169.		ing pairs does the first gas eduction while the second in [Manipal MEE 1995]  (b) $SO_2$ and $Cl_2$	,,,,,	sulphur when <i>X</i> is  (a) Fluorine or ch  (b) Chlorine only  (c) Chlorine and b	s llorine		[DCE 2003]
	(c) $H_2$ and $Br_2$	(d) $NH_3$ and $SO_2$		(d) $F$ , $Cl$ , $Br$ all	) 1 OHHIH	. 0111y	
170.	Which of the followin	g halogens does not form	40-			ough VCl	74T 1 74D

**180.** When  $I_2$  is passed through KCl, KF and KBr

[CPMT 2004]

solutions

(a)  $Cl_2$  and  $Br_2$  are evolved

(b)  $Cl_2$  is evolved

[MP PET 1997]

(b) Chlorine

(d) Iodine

oxyacid

(a) Fluorine

(c) Bromine

	798 s and p-Blo	ck Elements				
	(c) $Cl_2$ , $Br_2$ and $F_2$ are	evolved		(a) 5	(b) 1	
	(d) None of these			(c) 0	(d) None of these	
181.		increases in water in the	5.	Helium was discover		
	presence of			(a) Crooks	(b) Rutherford	
	1	[Pb. CET 2002]		(c) Frankland and Lo	ckyer (d)	Dorn
	(a) KI	(b) $H_2SO_4$	6.	The inert gases are	=	IT 1984]
	(c) $KMnO_4$	(d) $NH_3$		(a) Polyatomic	(b) Triatomic	
100	·	gen halides forms salts like		(c) Diatomic	(d) Monoatomic	
102.		alogen atom) [Kerala PMT 2004]	7•		ined at $100^{\circ}C$ absorbs	
	(a) <i>HF</i>	(b) <i>HCl</i>		(a) Ne and Kr	(b) He and Ar	
	(c) HI	(d) HBr	_	(c) Ar, Kr, Xe	(d) He and Ne	
	(e) All of these	(u) HBI	8.	Every inert gas atom		
182	With cold and dilute sodium hydroxide fluorine			(a) Has a saturated of		
103.	reacts to give [MH CET 2004]			<ul><li>(b) Has one electron</li><li>(c) Has eight electron</li></ul>		
	=	(b) $NaF + O_3$		(d) Has two electron		
	(c) $O_2$ and $O_3$		9.	Argon was discovered		MT 1991]
40.4	-	<del>-</del>	<i>y</i> .	(a) Rayleigh	(b) Frakland	and
184.	exhibit paramagnetic	ollowing oxides is expected	Lock			
	(a) $CO_2$	(b) $SO_2$		(c) Jansen	(d) Ramsay	
	<del>-</del>	_	10.	Deep sea divers used	to respirate is a mixto	ure of
_	(c) ClO <sub>2</sub>	(d) SiO <sub>2</sub>		(a) Oxygen and argor	n (b) Oxygen and he	elium
185.	Of the following acids	, the one that is strongest is		(c) Oxygen and nitro	gen (d) Oxygen	and
	(a) IID ()	[DPMT 2004]	-	rogen		
	(a) $HBrO_4$	(b) HOCl	11.	_	n forms maximum nui 1976; BHU 1980; CPMT	
	(c) $HNO_2$	(d) $H_3PO_3$		=	96; MP PMT 2001; Pb.CE	
186.	Which of the following is anhydride of perchloric			(a) <i>Ar</i>	(b) <i>He</i>	0.1
	acid			(c) <i>Xe</i>	(d) <i>Ne</i>	
	(-) Cl O	[CPMT 2004]	12.	Which of the fol	llowing gases exist	more
	(a) $Cl_2O_7$	(b) $Cl_2O_5$		abundantly in nature		
	(c) $Cl_2O_3$	(d) HClO			DPMT 1982,02; CPMT 1	
187.	$I_2$ dissolves in $KI$ solu	ition due to the formation of			anipal MEE 1995; MHCE	ET 2003]
		[CPMT 2004]		(a) Helium (c) Argon	(b) Neon	
	(a) $KI_2$ and $I^-$	(b) $K^+, I^-$ and $I_2$	13.	Which of the following	(d) Krypton	
	_	_	13.		RT 1976, 77; CPMT 1983,	86 001
	(c) $KI_3^-$	(d) None of these		(a) Nitrogen	(b) Fluorine	, 00, 90]
	Noble	daeoe		(c) Neon	(d) Oxygen	
	NODIE	gases	14.	Nuclear fusion produ		
_	TATIL'S C. Also C.	Name and the state of the state	•	(a) Argon	(b) Deuterium	
1.	Which of the fo	ollowing outer electronic nts argon[ <b>DPMT 1982; CPMT 197</b> 6	· NCE	•	(d) Krypton	
	configuration represe	Kurukshetra CEE 1998]	15.	Among the fluorides	below, the one whi	ch does
	(a) ns <sup>2</sup>	(b) $ns^2np^6$		not exist is	,	
					[NCERT 1977; CPM	IT 1988]
	(c) $ns^2np^5$	(d) $ns^2np^4$		(a) $XeF_4$	(b) $HeF_4$	
2.	Which mineral was us	sed in isolation of radium		(c) $SF_4$	(d) <i>CF</i> <sub>4</sub>	
	(a) Lime stone	[CPMT 1978, 81, 91]	16.	·	on would have electron	ıs
	(c) Rutile	<ul><li>(b) Pitch blende</li><li>(d) Haematite</li></ul>		3280		1971, 78]
3.	Which is the lightest a			(a) 6	(b) 2	- · · · · ·
ۍ.	(a) Hydrogen	(b) Oxygen		(c) 18	(d) 8	
	(c) Helium	(d) Nitrogen	17.	The electronic config		
4.	The valency of inert g	_	-	_	80, 81; DPMT 1982; MN	NR 1995]

				s and p-Block Elements 799
	(a) $1s^2$ , $2s^2 2p^2$	(b) $1s^2$ , $2s^2 2p^6$		(c) It is readily miscible with oxygen
	(c) $1s^2$ , $2s^2$			(d) It is less poisonous than nitrogen
18.		e tubes for advertisement	29.	Which of the following statements is not correct for a noble gas
		CPMT 1980, 89; MP PET 2002]		(a) Ar is used in electric bulbs
	(a) Argon	(b) Neon		(b) Kr is obtained during radioactive
	(c) Helium	(d) Xenon	disii	ntegration
19.	Least chemical activit	y is shown by [CPMT 1973, 79]		(c) Half life of <i>Rn</i> is only 3.8 days
	(a) Nitrogen	(b) Argon		(d) <i>He</i> is used in producing very low temperature
	(c) Methane	(d) Ammonia	30.	6 6 1
20.	Noble gases do not	react with other elements		a noble gas
	because			[CPMT 1976, 83, 89; BHU 1982; Pb. CET 2000 NCERT 1979; IIT Screening 1993; EAMCET 1993]
	( ) == 1	[CPMT 1981]		(a) $1s^2$ , $2s^22p^6$ , $3s^2$
	(a) They have complete electron shells	etely paired up and stable		(a) $1s$ , $2s$ $2p$ , $3s$ (b) $1s^2$ , $2s^22p^6$ , $3s^1$
	(b) The sizes of their	atoms are very small		· · · / • • /
	(c) Are not found in a	bundance		(c) $1s^2$ , $2s^2 2p^6$
	(d) Are monoatomic			(d) $1s^2$ , $2s^22p^6$ , $3s^23p^6$ , $4s^2$
21.	Monazite is source of		31.	Which of the following has zero valency[DPMT 1985]
	(a) He	(b) <i>Kr</i>	_	(a) Sodium (b) Beryllium
	(c) <i>Ar</i>	(d) <i>Ne</i>		(c) Aluminium (d) Krypton
22.		ring fluorides of Xenon is	32.	The forces acting between noble gas atoms are
	impossible	ıkshetra CEE 1998;RPET 1999]	•	[NCERT 1989]
	(a) $XeF_6$	(b) $XeF_4$		(a) Vander Waals forces
	(c) $XeF_3$	(d) $XeF_2$		(b) Ion-dipole forces
	-	(d) Aer 2		(c) London dispersion forces
23.	<i>XeF</i> <sub>2</sub> molecule is			(d) Magnetic forces
	(a) Square planer		33.	Which of the following is the correct sequence of
	(b) Trigonal bipyrami	dal		the noble gases in their group in the periodic
	(c) Trigonal planer			table
24.	(d) Linear	lysis produces [AFMC 1995]		[Manipal MEE 1995]
24.				(a) Ar, He, Kr, Ne, Rn, Xe(b) He, Ar, Ne, Kr, Xe, Rn
	(a) $XeF_2$	(b) XeOF <sub>2</sub>		(c) He, Ne, Ar, Kr, Xe, Rn(d) He, Ne, Kr, Ar, Xe, Rn
	(c) $XeOF_4$	(d) $XeO_3$	34.	Which of the following represent nobel gas configuration
25.	In $XeF_2$ hybridisation	of Xe is		[BHU 1995]
	(a) $sp^2$	(b) $sp^3d$		(a) $1s^2$ , $2s^22p^6$ , $3s^23p^63d^{10}$ , $4s^24p^64d^{10}$ , $5s^25p^6$
	(c) $sp^3$	(d) $sp^3d^2$		(b) $1s^2$ , $2s^22p^6$ , $3s^23p^63d^{10}$ , $4s^24p^64d^{10}4f^{14}$ ,
26.		llowing noble gases is the S 1983; MP PET 1999; Pb. PMT 20	001:	$5s^25p^65d^1,6s^2$
	F F	JIPMER (Med.) 2002]	<b>,</b>	(c) $1s^2$ , $2s^22p^6$ , $3s^23p^63d^{10}$ , $4s^24p^64d^{10}$ ,
	(a) <i>Xe</i>	(b) <i>Ar</i>		$5s^25p^65d^1$ , $6s^2$
	(c) Ne	(d) He		-
27.		llowing noble gases is not		(d) $1s^2$ , $2s^22p^6$ , $3s^23p^63d^{10}$ , $4s^24p^64d^{10}$
-	found in the atmosphe		35.	$XeF_6$ on hydrolysis gives
	(a) Rn	(b) <i>Kr</i>		[MP PET 2000; Pb. PMT 2000; DCE 2002]
	(c) <i>Ne</i>	(d) <i>Ar</i>		(a) $XeO_3$ (b) $XeO$

(c) *XeO*<sub>2</sub>

He, Ne, Ar, Kr, Xe, is

(a) He > Ne > Ar > Kr > Xe

36.

(d) Xe

[AIIMS 2002]

The correct order of solubility in water for

28. Helium is added to the oxygen supply used by

(b) It is lighter than nitrogen

pressure

deep sea divers because [MP PMT 1993; MP PET 1997]

(a) It is less soluble in blood than nitrogen at high

	800 s and p-Bloc	k Elements				
	(b) $Ne > Ar > Kr > He > 2$	Ke		48.		ng exhibits the weakest
	(c) $Xe > Kr > Ar > Ne > R$	<b>I</b> e			intermolecular forces	[KCET (Med.) 2001]
	(d) $Ar > Ne > He > Kr > X$	Ke			(a) He	(b) HCl
37.	In $XeF_2$ , $XeF_4$ , $XeF_6$ the	number of lo	one pairs on <i>Xe</i>		(c) $NH_3$	(d) $H_2O$
	is respectively		[AIEEE 2002]	49.	Which of the following a	re formed by Xenon
	(a) 2, 3, 1	(b) 1, 2, 3				[Roorkee 2000]
	(c) 4, 1, 2	(d) 3, 2, 1			(a) $XeF_3$	(b) <i>XeF</i> <sub>4</sub>
38.	Noble gases are group	of elements	which exhibit		(c) $XeF_5$	(d) <i>XeF</i> <sub>6</sub>
	very			50.	Among the following mo	lecule
	(a) IIi ah ahami al astin	_	la (Med.) 2002]		(i) $XeO_3$ (ii) $XeOF_4$ (iii)	
	(a) High chemical activ	=			•	mber of lone pairs on <i>Xe</i>
	(b) Low chemical activi	-		are	more naving bame na	moer or tone pairs on he
	<ul><li>(c) Minimum electrone</li><li>(d) Much paramagnetic</li></ul>	-				[AIIMS 2005]
20	Which noble gas is mos		water [CDMT 2002]		(a) (i) and (ii) only	(b) (i) and (iii) only
39.	(a) He	(b) <i>Ar</i>	vater [CFWI 2002]		(c) (ii) and (iii) only	(d) (i),(ii) and (iii)
	(c) Ne	(d) Xe		51.		ng first prepared a stable
40.	Gradual addition of ele	. ,	ls in the noble		compound of noble gas	[MP PET 1999]
1	gases causes a decrease		[MP PET 1997]		(a) Rutherford	(b) Rayleigh
	(a) Ionization energy	(b) Atomic	radius		(c) Ramsay	(d) Neil Bartlett
	(c) Boiling point	(d) Densit	y	52.		t gas elements is[MP PMT 1999]
41.	Which of the following				(a) Helium	(b) Neon
	octet of electrons in its				<ul><li>(c) Argon</li><li>Which of the following g</li></ul>	(d) Radon
	(a) Neon	(b) Radon		53.		[CPMT 2000; Pb. CET 2002]
	(c) Argon	(d) Helium			(a) Ne	(b) He
42.	The low chemical react be attributed to their	-	•		(c) Kr	(d) All of these
	(a) Being non-metals	ı	Pune CET 1998]	54.		ving statements regarding
	(b) Having high ionizat	ion energies			helium is incorrect	[AIEEE 2004]
	(c) Being gases	ion energies				ce and sustain powerful
	(d) Found in nature in s	small quanti	ties		superconducting mag	
43.	Percentage of $Ar$ in air	-	[CPMT 1989]		(b) It is used as a cryoge experiments at low t	enic agent for carrying out
	(a) 1%	(b) 2%			-	gas balloons instead of
	(c) 3%	(d) 4%				it is lighter and non-
44.	Which of the following	is not obta	ined by direct		inflammable	5
	reaction of constituent	elements	[MP PET 1994]		(d) It is used in gas-cool	ed nuclear reactors
	(a) $XeF_2$	(b) <i>XeF</i> <sub>4</sub>		55.	Which of the following is	nert gas liquifies easily
	(c) $XeO_3$	(d) <i>XeF</i> <sub>6</sub>				[Pb. CET 2002]
45.	Fluorine forms chemica	l compounds	s with[ <b>MP PMT 199</b> 4	4]	(a) Kr	(b) He
	(a) He	(b) <i>Ne</i>		_	(c) Ne	(d) <i>Ar</i>
	(c) <i>Ar</i>	(d) <i>Xe</i>		56.	The oxidation number of	Exenon in $XeOF_2$ is [J & K 2005]
46.	Which of the following	has $sp^3$ hyb	ridisation[DCE 200	1]	(a) Zero	(b) 2
	(a) <i>XeO</i> <sub>3</sub>	(b) <i>BCl</i> <sub>3</sub>			(c) 4	(d) 3
	(c) $XeF_4$	(d) $BBr_3$		57•		nighest boiling point[BCECE 200
4	•	-	and V C		(a) <i>Xe</i>	(b) Ar
47.	Which element out of			_ ^	(c) Kr	(d) He
	least number of compou	(b) <i>Ar</i>	[MP PMT 1995]	58.	=	s an inert gas [AFMC 2005]
	(c) Kr	(d) <i>Xe</i>			(a) $H_2$	(b) $O_2$
	(C) III	(u) Ae			(c) $N_2$	(d) Argon

**59.** Which of the following is most polarised[DPMT 2005]

- (a) *Kr*
- (b) *He*
- (c) Ar
- (d) Xe
- Which of the following is planar
- [J & K 2005]

- (a)  $XeF_2$
- (b)  $XeO_3F$
- (c)  $XeO_{2}F_{2}$
- (d)  $XeF_4$

## Critical Thinking Objective Questions

- 1. The correct sequence in decreasing order of the percentage of nitrogen in the given compounds is[NDA 1999]
  - (a) Urea > Ammonium chloride > Ammonium nitrate > Ammonium nitrite
  - (b) Urea > Ammonium nitrate > Ammonium nitrite > Ammonium chloride
  - (c) Urea > Ammonium nitrite > Ammonium nitrate > Ammonium chloride
  - (d) Urea > Ammonium nitrite > Ammonium chloride > Ammonium nitrate
- As the alkaline earth metals (except Be) tend to 2. lose their valence electrons readily they act as[Kerala (Med.)(2903]boxide
  - (a) Weak oxidising agent
- (b)Weak reducing agent
- (c) Strong oxidising agent (d)Strong reducing agent
- The first ionisation energies of alkaline earth 3. metals are higher than those of the alkali metals. This is because

### [UPSEAT 2001]

- (a) There is increases in the nuclear charge of the alkaline earth metals
- (b) There is decreases in the nuclear charge of the alkaline earth metals
- (c) There is no change in the nuclear charge
- (d) None of these
- Lead is maximum in 4.

[BVP 2004]

- (a) Soda glass (c) Pyrex glass
- (b) Jena glass (d) Flint glass
- 5.  $BaSO_4$  and carbon on heating reacts to produce

### [Pb. PMT 2004]

- (a)  $Ba + SO_2 + CO_2$
- (b) BaS + CO
- (c)  $BaS + O_2 + SO_3$
- (d)  $BaCO_3 + S + O_2$
- 6. The atomic radii of alkali metals (M) lie in the order Li < Na < K < Rb but the radii of  $M^+$  ions in aqueous solution lie in the reverse order  $Li^+ > Na^+ > K^+ > Rb^+$ . What is the reason for this reverse order (on going from Li to Rb)?[MP PMT 1997]
  - (a) Gradual increase in ionisation energy
  - (b) Increasing weakness of the metallic bond
  - (c) Increasing electropositive character

- (d) Decreasing degree of hydration
- Fusion mixture is 7.

[CPMT 2002]

- (a)  $Na_2CO_3 + K_2CO_3$
- (b)  $Na_2CO_3 + NaHCO_3$
- (c)  $Na_2CO_2 + NaOH$
- (d)  $Na_2CO_3 + K_2SO_4$
- 8. Concentrated hydrochloric acid when kept in open air sometimes produces a cloud of white fumes. The explanation for it is that
  - (a) Concentrated hydrochloric acid emits strongly smelling HCl gas all the time
  - (b) Oxygen in air reacts with the emitted HCl gas to form a cloud of chlorine gas
  - (c) Strong affinity of HCl gas for moisture in air results in forming of droplets of liquid solution which appears like a cloudy smoke
  - (d) Due to strong affinity for water, concentrated hydrochloric acid pulls moisture of air towards itself. This moisture forms droplets of water and hence the cloud
- Match List I with List II and select the correct answer using the codes given below the lists

#### List I

(a) Peroxide

- List II (1)  $C_3O_2$
- (b) Superoxide
- (2)  $PbO_2$
- (c) Dioxide
- (3) KO<sub>2</sub>
- (4)  $H_2O_2$

Codes:

(a) A В C D 3 2 4 1

- (b) A В C D
  - 2 1 3 4
- C (c) A
- 4 2 3 1 (d) A C D R
  - 2 3
- The most efficient agent for the absorption of  $SO_3$  is

[KCET 1998]

[NDA 1999]

- (a)  $98\%H_2SO_4$
- (b)  $80\%H_2SO_4$
- (c) 20% oleum
- (d)  $90\% H_2 SO_4$
- Mark the oxide which is amphoteric in character 11. [MP PMT 2000]
  - (a)  $CO_2$
- (b) SiO<sub>2</sub>
- (c)  $SnO_{2}$
- (d) CaO
- Concentrated aqueous sodium hydroxide can separate mixture of [MNR 1995]
  - (a)  $Al^{3+}$  and  $Sn^{2+}$
- (b)  $Al^{3+}$  and  $Fe^{3+}$
- (c)  $Al^{3+}$  and  $Zn^{2+}$
- (d)  $Zn^{2+}$  and  $Pb^{2+}$
- The composition of the common glass is[DCE 2004] (a)  $Na_2O.CaO.6SiO_3$ 
  - (b)  $Na_2O.Al_2O_3.SiO_2$
  - (c)  $CaO.Al_2O_3.SiO_2$
- (d)  $Na_2O.CaO.6SiO_2$

14.		not form ammonium nitrate		(c) Sea shells	(d) Dolomit	
	by reaction with dilute		25.[]		llowing statements	is false for
	(a) <i>Al</i>	(b) Fe		alkali metals	FMAID 4004	MD DET accel
1-	(c) Pb	(d) Mg		(a) Lithium is the	strongest reducing	MP PET 2001]
15.	rotal number of lone p	pair of electrons in $XeOF_4$ is		(b) <i>Na</i> is amphot		agent
	(-) 0	[IIT-JEE (Screening) 2004]		=		
	(a) 0	(b) 1		(c) $Li^+$ is exception		in in linuid
16.	(c) 2 What is the correct re	(d) 3 elationship between the <i>pHs</i>		ammonia	als give blue solut	non in iiquia
10.		s of sodium oxide $(pH_1)$ ,	26.		ne in water is grea	tly increased
		, sodium selenide $(pH_3)$ and			of iodide ions be	5
	sodium telluride $(pH_4)$	- 5		formation of		
	(F4)	[CBSE PMT 2005]				[IIT 1994]
	(a) $pH_1 > pH_2 = pH_3 > p$			(a) $I_2$	(b) $I_3$	
	(b) $pH_1 < pH_2 < pH_3 < pH_4 < pH_5 < pH_$	·		(c) $I_3^-$	(d) $I^-$	
	(c) $pH_1 < pH_2 < pH_3 = p$	·	27.	The solubility in v	water of sulphates	down the Be
		·	,	_	>> Ca > Sr > Ba . This	
	(d) $pH_1 > pH_2 > pH_3 > p$	·			[CI	BSE PMT 1995]
17.		owing is not an amphoteric		(a) High heat of	solvation for small	ller ions like
	substance	[KCET 2004]	$Be^{2+}$			
	(a) $HNO_3$	(b) HCO <sub>3</sub>		(b) Increasing mo	lecular weight	
	(c) $H_2O$	(d) $NH_3$		(c) Decreasing lat		
-0	-			(d) Increase in me	~ .	
18.	table	buffer group of the periodic	28.	Magnesium burns	•	
	tubic	[Pb. CET 2004]		(-) 1(-0	[CPMT 1988, 8	9; AFMC 1987]
	(a) I	(b) VII		(a) MgO	(b) $Mg_3N_2$	_
	(c) VIII	(d) Zero		(c) $MgCO_3$	(d) <i>MgO</i>	and $Mg_3N_2$
19.	Which of the following	g salt is insoluble in water	both			
		[MP PET 2004]	29.	-	ol when heated w	
	(a) $CuSO_4$	(b) $CdSO_4$		=	=	ne compound[ <b>CPMT 1</b>
	(c) $PbSO_4$	(d) $Bi_2(SO_4)_3$		(a) $BaZnO_2$	(b) $Ba + ZnC$	_
20.	Which of the following	goxides is the most acidic		(c) $BaCdO_2$	(d) $BaO_2 + 2$	Zn
		BSE PMT 1999; MP PMT 2002]	30.	Which of the follo	wing oxide is diama	agnetic
	(a) $N_2O_5$	(b) $P_2O_5$				[MP PET 1990]
	(c) $As_2O_5$	(d) $Sb_2O_5$		(a) <i>NO</i>	(b) $N_2O_4$	
21.	Whose bond energy is			(c) $NO_2$	(d) $N_2O_5$	
	(a) $F_2$	[CPMT 1988; MP PMT 1990] (b) Cl <sub>2</sub>	31.		lowing salt becom propriately hydrate	-
	(c) $Br_2$	(d) $I_2$		(a) $ZnCO_3$	(b) CaSO <sub>4</sub>	a [CIMI 1905]
22.		reatment with steam under		3	•	
22.	pressure gives $NH_3$ ar			(c) MgSO <sub>4</sub>	(d) CaCO <sub>3</sub>	
	(a) CaHCO <sub>3</sub>	(b) CaO	32.	alkaline earth met	lectron and proton al ion will be	in the third
	(c) $Ca(OH)_2$	(d) CaCO <sub>3</sub>		(a) $\frac{e}{20}, \frac{p}{20}$	(b) $\frac{e}{18}, \frac{p}{20}$	
23.	Six volumes of oxygen	n, on complete ozonisation,		20, 20	18, 20	
	form volumes of o	=	[]	DPMT <u><b>20</b>0<b>0</b>]</u> 18, 18	(d) $\frac{e}{19}, \frac{p}{20}$	
	(a) 4	(p) 3		18 18	19 20	
	(c) 2	(d) 6	33.	_	f alkaline earth me	tals have the
24.	The substance not like	ly to contain $CaCO_3$ is		following magneti		HDMED 2022
		[AIEEE 2003]		(a) Diamagnetic	T 1998; RPMT 2000; (b) Parama	
	(a) A marble statue	(b) Calcined gypsum		(a) Diamagnetic	(U) Farailia	5110410

s and p-Block Elements 803  $KO_2$  (potassium superoxide) is used in oxygen (c) Ferromagnetic (d) Diaferromagnetic 42. Which of the following is the life saving mixture cylinders in space and submarines because it [AIEEE 2002 34. for an asthma patient [MP PMT 2001] (a) Absorbs  $CO_2$  and increases  $O_2$  content (a) Mixture of helium and oxygen (b) Eliminates moisture (b) Mixture of neon and oxygen (c) Absorbs  $CO_2$ (c) Mixture of xenon and nitrogen (d) Produces ozone (d) Mixture of argon and oxygen Which would quickly absorbs oxygen Fire extinguishers contain  $H_2SO_4$  and [AFMC 1980] [CBSE PMT 1992; MP PET 1995] (b)  $Na_2CO_3$ (a) CaCO<sub>3</sub> (a) Alkaline solution of pyrogallol (d)  $NaHCO_3$  and  $Na_2CO_3$ (c) NaHCO<sub>3</sub> (b) Conc.  $H_2SO_4$ Which is insoluble in water (c) Lime water [CPMT 2003] (d) Alkaline solution of CuSO A (a)  $H_2S$ (b)  $HgCl_2$ Nitrogen is liberated the thermal (c)  $Ca(NO_3)_2$ (d)  $CaF_2$ decomposition of only Which of the following halides is most acidic[KCET 1996] 45. [IIT 1991] (a)  $PCl_3$ (b) SbCl<sub>3</sub> (a)  $NH_4NO_2$ (b)  $NaN_3$ (c)  $BiCl_3$ (d) CCl<sub>4</sub> (c)  $(NH_A)_2 Cr_2 O_7$ (d) All the three The stability of the following alkali metal 46. 37. Red phosphorus is less reactive than yellow chlorides follows the order phosphorus because (a) LiCl > KCl > NaCl > CsCl[DPMT 1982; JIPMER 1999; CBSE PMT 1999; RPET 2003] (b) CsCl > KCl > NaCl > LiCl(a) Its colour is red (c) NaCl > KCl > LiCl > CsCl(b) It is highly polymerised (d) KCl > CsCl > NaCl > LiCl(c) It is hard 47. The reaction of  $Na_2S_2O_3$  with iodine gives (d) It is insoluble in  $C_2H_5OH$ [CPMT 1971, 80, 81; DPMT 1983, 90; Carbon differs from other elements of the group. MP PMT 1985; EAMCET 1990; BHU 1980] Which is the false statement (a) Sodium sulphide (b) Sodium sulphite (a) Due to its marked tendency to form long (c) Sodium sulphate (d) Sodium tetrathionate chains (catenation) Which one of the following is the true covalent 48. (b) Due to its unique ability to form multiple oxide of iodine [MP PET/PMT 1988] (a)  $I_2O_4$ (b)  $I_2O_5$ (c) Due to *d*-orbital in penultimate shell (c)  $I_2O_7$ (d)  $I_2O_9$ (d) Due to its limitation of co-ordination number Lithium aluminium hydride acts as 49. [CPMT 1994] (a) Oxidising agent Which of the following oxide does not form acidic (b) Reducing agent 39. aqueous solution [CPMT 2004] (c) Both the above (d) None of these (a)  $N_2O_3$ (b)  $NO_2$ The mixture of conc. HCl and potassium chlorate on heating gives [Roorkee 2000] (c)  $N_2O_5$ (d) NO (a)  $Cl_2$  only (b) ClO2 only 40. Which of the following is in the increasing order (c)  $Cl_2 + ClO_2$ (d)  $Cl_2 + ClO_2 + ClO_3$ of the ionic character [JIPMER 2002] When SO, is passed through acidified solution of (a)  $PbCl_4 < PbCl_2 < CaCl_2 < NaCl$ 51. (b)  $PbCl_2 < PbCl_4 < CaCl_2 < NaCl$ [CPMT 1973, 81, 93] (c)  $PbCl_2 < PbCl_4 < NaCl < CaCl_7$ (a)  $H_2SO_4$  is formed (b)  $H_2SO_3$  is formed (d)  $PbCl_4 < PbCl_2 < NaCl < CaCl_2$ (c) Sulphur is precipitated (d) None of these Silicon chloroform is prepared by 41. [MH CET 1999] Four reactions are given below (a) Si + HCl(b)  $SiCl_4 + H_2O$ (i)  $2Li + 2H_2O \rightarrow 2LiOH + H_2$ (c)  $SiF_A + NaF$ (d)  $H_2SiF_6 + Cl_2$ 

(ii)  $2Na + 2H_2O \rightarrow 2NaOH + H_2$ (iii)  $2LiNO_3 \xrightarrow{Heat} 2LiNO_2 + O_2$ 

(iv)  $2NaNO_3 \xrightarrow{Heat} 2NaNO_2 + O_2$ Which of the above, if any, is wrong (a) (iv) (b) (iii) (c) (i) (d) None of these

53. Increasing order of solubility is [AFMC 1987]

(a) CaCO<sub>3</sub>, KHCO<sub>3</sub>, NaHCO<sub>3</sub>

(b) NaHCO<sub>3</sub>, KHCO<sub>3</sub>, CaCO<sub>3</sub>

(c) KHCO<sub>3</sub>, NaHCO<sub>3</sub>, CaCO<sub>3</sub>

(d) CaCO<sub>3</sub>, NaHCO<sub>3</sub>, KHCO<sub>3</sub>

Nitrolim is [CPMT 1976, 78, 2000; BHU 1987; DCE 1999; RPMT 2000]

(a)  $Ca(NO_3)_2$ 

(b)  $Ca(CN)_2$ 

(c)  $CaCN_2 + C$ 

(d)  $CaCN_2$ 

55. The following acids have been arranged in the order of decreasing acid strength. Identify the correct order.

(I) ClOH (II) BrOH (III) IOH

(a) I > II > III

(b) II > I > III

(c) III > II > I

(d) I > III > II

Which of the following element does not belong to the family of elements indicated [Orissa JEE 1997]

(a) Rubidium (Rb, Z = 37): Alkali metals

(b) Barium (Ba, Z = 56): Alkaline earth metals

(c) Iridium (I, Z = 77): Nobel gases

(d) Argon (Ar, Z = 18): Nobel gases

57.  $H_3PO_2$  is the formula for one of the phosphorus acid. Its name and basicity are respectively

[CBSE PMT 1992; BHU 1999; KCET 1999]

(a) Phosphorus acid and two

(b) Hypophosphorus acid and two

(c) Hypophosphorus acid and one

(d) Hypophosphoric acid and two

58. Which of the following oxides of nitrogen is paramagnetic

[CPMT 1984; CBSE PMT 1994; AIIMS 2000]

(a)  $N_2O_3$ 

(b)  $N_2O$ 

(c)  $NO_2$ 

(d)  $N_2O_5$ 

59. Nessler's reagent is

[CPMT 2002]

(a) Potassium in mercuric iodide

(b)  $TiCl_{4}$ 

(c) Anhydrous AlCl<sub>3</sub>

(d)  $Al_2O_3 / Cr_2O_3$ 

**60.** The noble gas was first time discovered by

(a) Cavandish

(b) William Ramsay

(c) Rayleigh

(d) Frankland

The ratio of  $\frac{C_p}{C_v}$  for inert gases is [DCE 1999]

(a) 1.99

(b) 2.13

(c) 1.66

(d) 1.33

62. White P reacts with caustic soda, the products are  $PH_3$  and  $NaH_2PO_2$ . This reaction is an example of [DCE 20]

(a) Oxidation

(b) Reduction

(c) Disproportionation (d) Neutralisation

The oxide, which cannot act as a reducing agent, 63.

[Pb. CET 2002]

(a)  $NO_2$ 

(b) SO,

(c) CO<sub>2</sub>

(d) ClO<sub>2</sub>

Which of the following product is formed when SiF<sub>4</sub> reacts with water [Pb. CET 2003]

(a)  $SiF_3$ 

(b)  $H_4SiO_4$ 

(c)  $H_2SO_4$ 

(d)  $H_2SiF_4$ 

**65.** Ozone with dry iodine give

[Pb. CET 2003]

(a)  $I_4O_4$ 

(b)  $I_2O_3$ 

[IIT(a)9g6]

(d)  $I_2O_4$ 

66. The microcosmic salt is[Pb.CET 2004; Pb. PMT 2004]

(a)  $Na(NH_A)H_2O$ 

(b)  $K(NH_4)HPO_32H_2O$ 

(c)  $Na(NH_4)HPO_44H_2O$ ) (d)  $Na(NH_3)HPO_44H_2O$ 

Thermite is a mixture of

[Pb. CET 2004]

(a)  $Cr_2O_3 + Al_2O_3$ 

(b)  $Fe_2O_3 + Al$ 

(c)  $Fe_2O_3 + Al_2O_3$ 

(d)  $Al_2O_3 + 2Cr$ 

**68.** The colour of liquid  $O_2$  is

[BVP 2004]

(a) Red

(b) Dark blue

(c) Pale yellow

(d) Pale blue

Which of the following gas mixture is used by the divers inside the sea [AFMC 2004]

(a)  $O_2 + He$ 

(b)  $O_2 + Xe$ 

(c)  $O_2 + Ar$ 

(d)  $O_2 + N_2$ 

One mole of magnesium nitride on the reaction 70. with an excess of water gives

(a) Two moles of ammonia (b)One mole of nitric acid

(c) One mole of ammonia

(d)Two moles of nitric acid

Calcium cyanamide on treatment with steam produce

[Pb. PMT 2004]

(a)  $CaCO_3 + NH_3$ 

(b)  $CaHCO_3 + NH_3$ 

(c)  $CaO + NH_3$ 

(d)  $Ca(OH)_2 + NH_3$ 



Read the assertion and reason carefully to mark the correct option out of the options given below:

(a)	_	ertion and reason are true and the		Reason :	The catenated $-O-O-O-$ changes are less stable as compared to
(b)		correct explanation of the assertion. tion and reason are true but reason is			O = O molecule. [AIIMS 2001]
(c)	not the corre	ct explanation of the assertion. s true but reason is false.	10.	Assertion:	Potassium and caesium are used in photo-electric cells.
(d)		on and reason both are false.		Reason :	Potassium and caesium emit
(e)	•	s false but reason is true.			electrons on exposure to light.[AIIMS 2002]
			11.	Assertion :	The fluorine has lower reactivity.
1.	Assertion :	Sulphate is estimated as $BaSO_4$ and		Reason :	F-F bond has low bond
		not as $MgSO_4$ .	12.	Assertion:	dissociation energy. [AIIMS 2002] Halogens do not occur in free state.
	Reason :	Ionic radius of $Mg^{2+}$ is smaller than	12,	Reason :	Halogens are highly reactive. [AIIMS 1994]
		that of $Ba^{2+}$ . [IIT 1998]	13.	Assertion :	Lithium forms Lithium oxide $(LiO_2)$ .
2.	Assertion:	Amongst the halogens fluorine can	_5.	Reason :	$N_2$ molecule have unpaired
		oxidise the elements to highest oxidation state.		Reason .	electrons.
	Reason :	Due to small size of fluoride ion, it	1.4	Assertion:	[AIIMS 1995] Liquid $NH_2$ is used for
		is difficult to oxidise fluoride ion to fluorine. Hence reverse reaction	14.	Assertion.	Liquid $NH_3$ is used for refrigeration.
		takes place more easily. [IIT 1996]		Reason :	Liquid <i>NH</i> <sub>3</sub> quickly vaporises.
3.	Assertion:	$HNO_3$ is a stronger acid than		Reason .	[AIIMS 1995]
		$HNO_2$ .	15.	Assertion:	$Al(OH)_3$ is insoluble in $NH_4OH$ but
	Reason :	In $\mathit{HNO}_3$ there are two nitrogen-to-	-3.	11000101011	soluble in NaOH.
		oxygen bonds whereas in $\mathit{HNO}_2$		Reason :	NaOH is strong alkali. [AIIMS 1997]
		there is only one.	16.	Assertion :	Boron is metalloid.
4.	Assertion :	The Value of Vander Waal's		Reason :	Boron shows metallic nature. [AIIMS 1997]
		constant 'a' is larger for ammonia than for nitrogen.	17.	Assertion:	Inert gases are monoatomic.
	Reason :	Hydrogen bonding is present in ammonia.		Reason :	Inert gases have stable configuration.
		[IIT 1998]			[AIIMS 1999]
5.	Assertion:	Xenon forms fluorides.	18.	Assertion :	Magnesium continue to burn in nitric oxide.
	Reason :	Due to the strong electronegativity of fluorine . [AIIMS 2001]		Reason :	During burning heat evolved do not
6.	Assertion:	Chlorine and sulphur dioxide both	19.	Assertion:	decompose $NO$ . [AIIMS 2001] Anhydrous $BaO_2$ is used for
	Reason :	are bleaching agents.  The bleaching action of chlorine	19.	Assertion.	preparing $H_2O_2$ .
	Reason .	and sulphur dioxide is performed		Reason :	Hydrated $BaO_2$ is not available.
		through the process of oxidation.[AIIMS	2000	]	[AIIMS 2001]
7•	Assertion:	Nitrogen is unreactive at room	20.	Assertion:	Benzene is reactive while inorganic
		temperatures but becomes reactive at elevated temperatures (On			benzene is unreactive compound.
		heating) or in presence of catalysts.		Reason :	Inorganic benzene is,
	Reason :	In nitrogen molecule, there is			borazine, $B_3N_3H_6$ .
		extensive delocalization of			[AIIMS 2002]
•	A	electrons.	21.	Assertion:	Halogens absorb visible light.
8.	Assertion:	Covalency of oxygen is three.		Reason :	All halogens are coloured.[AIIMS 2002]
	Reason :	Dinegative anion of oxygen $(O^{2-})$ is quite common but dinegative anion		Assertion:	Barium is not required for normal biological function in human.
		of sulphur $(S^{2-})$ is less common.[AIIMS	2001]	Reason :	Barium does not show variable
9.	Assertion:	At room temperature, oxygen exists as a diatomic gas, whereas sulphur		A 000-+	oxidation state. [AIIMS 2003]
		exists as solid.	23.	Assertion :	The $O-O$ bond length in $H_2O_2$ is
					shorter than that of $O_2F_2$ .
				Reason :	$H_2O_2$ is an ionic compound.[AIIMS 2003]

	806 s a	nd p-Block Elements			
24.	Assertion :	$PbI_4$ is a stable compound.	36.	Assertion :	Although $PF_5$ , $PCl_5$ and $PBr_3$ are
	Reason :	Iodide stabilizes higher oxidation state.			known, the pentahalides of nitrogen have not been observed.
25.	Assertion :	[AIIMS 2003]  Mg is not present in enamel of human		Reason :	Phosphorus has lower electronegativity than nitrogen.
		teeth.	37.	Assertion :	The electronic structure of $O_3$ is
	Reason :	<i>Mg</i> is an essential element for biological functions of human. [AIIMS :	2004]		
26.	Assertion :	Radium is most abundant <i>s</i> -block element.		December	
	Reason :	s-block elements are non-radioactive in nature.		Reason :	:0
27.	Assertion :	<i>LiCl</i> is predominantly a covalent compound.			Structure is not allowed because octet around <i>O</i> cannot be expanded.
	Reason :	Electronegativity difference between $Li$ and $Cl$ is too small.	38.	Assertion:	Sulphuric acid is more viscous than water.
28.	Assertion :	The first ionization energy of $Be$ is greater than that of $B$ .		Reason :	Concentrated sulphuric acid has a great affinity for water.
	Reason :	2 <i>p</i> -orbital is lower in energy than 2 <i>s</i> -orbital.	39.	Assertion:	$PCl_5$ is covalent in gaseous and
29.	Assertion :	The alkali metals can form ionic hydrides which contains the hydride ion.		Reason :	liquid states but ionic in solid state. $PCl_5$ in solid state consists of tetrahedral $PCl_4^+$ cation and
	Reason :	The alkali metals have low			octahedral $PCl_6^-$ anion.
		electronegativity, their hydrides conduct electricity when fused and liberate hydrogen at the anode.	40.	Assertion:	Among nitrogen halides $NX_3$ , the dipole moment is highest for
30.	Assertion :	Be does not impart any characteristic colour to the bunsen		Reason :	$NI_3$ and lowest for $NF_3$ . Nitrogen halides $NX_3$ , have trigonal
	Doncon	flame.			pyramidal structure.
	Reason :	Due to its very high ionization energy, beryllium requires a large amount of energy for excitation of	41.	Assertion :	White phosphorus is stored under water.
31.	Assertion :	the electrons.  Potassium is not obtained by the		Reason :	White phosphorous is highly reactive and catches fire
_		electrolysis of fused KCl.			spontaneously in air.
	Reason :	Potassium vapourises at the melting point of <i>KCl</i> .	42.	Assertion :	$Al$ forms $[AlF_6]^{3-}$ but $B$ does not form $[BF_6]^{3-}$ .
32.	Assertion:	Helium and beryllium have similar outer electronic configuration of		Reason :	$B$ does not react with $F_2$ .
	_	the type $ns^2$ .	43.	Assertion :	$NO_3^-$ is planar while $NH_3$ is
	Reason :	Both are chemically inert.		_	pyramidal.
33.	Assertion :	$Na_2SO_4$ is soluble while $BaSO_4$ is		Reason :	$N$ in $NO_3^-$ is $sp^2$ hybridized but in
	Dongon .	insoluble.			$NH_3$ it is $sp^3$ – hybridized.
	Reason :	Lattice energy of <i>BaSO</i> <sub>4</sub> exceeds its hydration energy.	44.	Assertion :	Si - Si bonds are much stronger then $Si - O$ bonds.
34.	Assertion :	Alkali metals impart colour to the flame.		Reason :	Silicon forms double bonds with itself.
	Reason :	Their ionisation energies are low.	45.	Assertion:	The $S-S-S$ bond angle in
35.	Assertion:	Superoxides of alkali metals are paramagnetic.		Reason :	$S_8$ molecule is 105°. $S_8$ has a $V$ -shape.
	Reason :	Superoxides contain the ion $O^{-2}$ which has one unpaired electron.			

**46.** Assertion: Caro's acid has S atom in +6

oxidation state.

Reason : Caro's acid contains one peroxo  $O_2^{2-}$ 

group.

47. Assertion: The m.p./b.p. of noble gases are

quite high.

Reason : The interparticle forces among

noble gases in their liquid state are

covalent forces.

**48.** Assertion: In  $SO_2$ , the bond angle is 119°

whereas in  $SO_3$ , the bond angle is

120°.

Reason: S atom in both  $SO_2$  and  $SO_3$  is

 $sp^2$  – hybridized.

**49.** Assertion: Calcium carbide on hydrolysis gives

methane.

Reason : Calcium carbide contains

 $C^{4-}$  anions.

50. Assertion: Xenon forms fluorides.

Reason : Because 5 d-obitals are available for

valence shell expansion.

51. Assertion: Hydrogen cannot be prepared in

laboratory.

Reason : Hydrogen of high purity is obtained

by electrolysing warm aqueous barium hydroxide between nickel

electrodes.

**52.** Assertion: Diprotium shows relatively inert

behaviour at room temperature.

Reason: The values of melting point and

boiling point for dideuterium are higher as compared to diprotium.

Water can be transformed from

**53.** Assertion: Water can be transform liquid to solid state only.

Reason : The distribution of water over the

earth surface is uniform.

**54.** Assertion: Ice is less dense than water.

Reason : Ice is a solid whereas water is

liquid.

**55.** Assertion:  $HF, NH_3$  and  $H_2O$  form

intermolecular hydrogen bonding.

Reason :  $HF, NH_3$  and  $H_2O$  molecules are

bonded in same manner.

56. Assertion: Hard water does not lather with

soap.

 $Reason \quad : \quad In \ hard \ water, \ the \ sodium \ stearate$ 

of soap changes to the corresponding calcium magnesium

salt which precipitates out.

**57.** Assertion :  $H_2O_2$  is stored in wax-lined glass.

Reason : Presence of metal surfaces, traces of alkali (present in glass) etc.

increases its decomposition.

**58.** Assertion:

A nearly tetrahedral arrangement of the orbitals about the oxygen atom allows each water molecule to form hydrogen bonds with as many as

four neighbouring water molecules.

Reason : In ice each water molecule form four hydrogen bond as each

four hydrogen bond as each molecule is fixed in the space.

**59.** Assertion: Calgon is used for removing  $Ca^{2+}$ 

and  $Mg^{2+}$  ions from hard water.

Reason : Calgon forms precipitate with

 $Ca^{2+}$  and  $Mg^{2+}$  ions.

**60.** Assertion: Reaction of  $SO_2$  and  $H_2S$  in the

presence of  $Fe_2O_3$  catalyst gives

elemental sulphur.

Reason :  $SO_2$  is a reducing agent.[AIIMS 2005]

**61.** Assertion :  $SiF_6^{2-}$  is known but  $SiCl_6^{2-}$  is not.

Reason : Size of fluorine is small and its lone

pair of electrons interacts with d-orbitals of Si strongly. [AIIMS 2005]

62. Assertion: Borax bead test is not suitable for

Al(III).

Reason :  $Al_2O_3$  is insoluble in water.[AIIMS 2005]

**63.** Assertion :  $SeCl_4$ , does not have a tetrahedral

structure.

Reason : Se in  $SeCl_4$  has two lone pairs.

[AIIMS 2005]

**64.** Assertion: Ozone is a powerful oxidizing agent

in comparison to  $O_2$ .

Reason: Ozone is diamagnetic but  $O_2$  is

paramagnetic. [AIIMS 2005]



#### Alkali metals b 2 С 3 4 b 5 а b 7 8 9 С а С 10 d 11 С 12 b 13 14 d 15 а 16 h 17 18 19 h 20 С b b 21 а 22 b 23 24 а 25 d 26 27 28 29 30 h h d С C 34 d 32 33 h 35 d а а 39 d 40 b 36 b 37 а 38 41 d 42 43 44 d 45 d c h

46	b	47	b	48	b	49	С	50	С
51	d	52	С	53	d	54	d	55	а
56	С	57	а	58	С	59	d	60	а
61	b	62	b	63	а	64	а	65	С
66	b	67	d	68	а	69	d	70	d
71	С	72	а	73	а	74	С	75	b
76	d	77	С	78	а	79	b	80	а
81	d	82	а	83	С	84	b	85	С
86	С	87	d	88	b	89	С	90	а
91	С	92	а	93	С	94	а	95	С
96	а	97	а	98	b	99	d	100	С
101	а	102	d	103	b	104	С	105	С
106	а	107	С	108	b	109	b	110	d
111	d	112	С	113	а	114	С	115	а
116	a	117	С	118	С	119	b	120	а
121	b	122	С	123	С	124	С	125	d
126	С	127	а	128	b	129	a,b	130	b
131	d	132	b	133	d	134	d	135	b
136	d	137	d	138	а	139	а	140	b
141	С	142	С	143	а	144	а	145	b
146	d	147	b	148	d	149	С	150	b

### Alkaline earth metals

1	С	2	d	3	а	4	b	5	b
6	d	7	d	8	d	9	b	10	a
11	а	12	d	13	а	14	а	15	а
16	d	17	С	18	С	19	b	20	b
21	b	22	d	23	d	24	а	25	а
26	bc	27	d	28	d	29	b	30	b
31	d	32	d	33	b	34	b	35	b
36	d	37	С	38	d	39	d	40	а
41	С	42	а	43	а	44	а	45	а
46	С	47	b	48	d	49	С	50	С
51	а	52	С	53	b	54	а	55	b
56	а	57	d	58	а	59	b	60	b
61	d	62	b	63	d	64	а	65	а
66	d	67	а	68	С	69	d	70	а
71	С	72	а	73	b	74	С	75	b
76	d	77	b	78	d	79	С	80	С
81	d	82	а	83	С	84	а	85	С
86	d	87	С	88	d	89	а	90	а
91	b	92	С	93	С	94	b	95	а
96	d	97	d	98	d	99	а	100	b

101	b	102	С	103	а	104	b	105	С
106	а	107	а	108	С	109	d	110	d
111	b								

## Boron family

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

### Carbon family

1	С	2	d	3	d	4	С	5	а
6	а	7	b	8	d	9	С	10	d
11	С	12	b	13	d	14	b	15	С
16	а	17	b	18	b	19	d	20	d
21	С	22	С	23	а	24	а	25	С
26	а	27	С	28	а	29	а	30	d
31	а	32	а	33	d	34	b	35	b
36	b	37	С	38	b	39	b	40	b
41	а	42	С	43	b	44	С	45	d
46	С	47	С	48	b	49	d	50	а
51	С	52	С	53	а	54	а	55	а
56	С	57	b	58	b	59	а	60	b
61	d	62	а	63	d	64	С	65	d
66	b	67	С	68	а	69	b		

### Nitrogen family

1	b	2	а	3	b	4	b	5	а
6	а	7	d	8	b	9	а	10	b
11	ad	12	b	13	а	14	b	15	d

16	b	17	d	18	b	19	С	20	а
21	а	22	а	23	b	24	d	25	С
26	С	27	С	28	С	29	d	30	С
31	d	32	С	33	d	34	ab	35	а
36	а	37	b	38	d	39	b	40	С
41	b	42	а	43	b	44	d	45	b
46	d	47	d	48	а	49	b	50	b
51	а	52	d	53	С	54	d	55	d
56	С	57	b	58	С	59	С	60	С
61	а	62	d	63	b	64	а	65	b
66	С	67	С	68	а	69	а	70	b
71	а	72	а	73	а	74	b	75	а
76	b	77	С	78	b	79	С	80	d
81	d	82	d	83	d	84	С	85	d
86	d	87	С	88	b	89	d	90	С
91	а	92	b	93	d	94	b	95	d
96	С	97	С	98	b	99	а	100	d
101	d	102	b	103	d	104	а	105	d
106	d	107	d	108	а	109	d	110	а
111	а	112	С	113	d	114	С	115	а
116	а	117	С	118	b	119	а	120	С
121	b	122	b	123	d	124	d	125	С
126	b	127	b	128	С	129	d	130	b
131	С	132	b	133	b	134	b	135	b
136	а	137	а	138	d	139	а	140	d
141	а	142	С	143	b	144	а	145	а
146	d	147	b	148	b	149	а	150	d
151	d	152	d	153	а	154	е	155	а
156	а	157	С	158	d	159	b	160	d
161	С	162	b	163	С	164	b	165	d
166	а	167	а	168	а	169	b	170	d
171	С	172	d	173	а	174	d	175	С
176	b	177	d	178	С	179	b	180	а
181	b	182	а	183	d	184	С	185	а
186	d	187	d	188	а	189	а	190	d
191	а	192	а	193	С	194	а	195	b
196	С	197	b	198	а	199	b	200	С
201	d	202	С	203	b	204	а	205	b
206	а	207	С	208	d	209	С	210	С
211	а	212	С	213	d	214	b	215	b,c
216	b	217	а	218	d	219	d	220	С
221	а	222	а	223	а	224	b	225	а
226	d	227	b	228	b	229	b	230	С

231	a	232	а	233	С	234	а	235	d
236	а	237	b	238	d	239	b	240	С
241	а								

## Oxygen family

1	b	2	С	3	b	4	С	5	а
6	С	7	b	8	d	9	b	10	а
11	а	12	b	13	bc	14	d	15	d
16	а	17	b	18	С	19	а	20	а
21	а	22	d	23	а	24	С	25	С
26	d	27	d	28	b	29	b	30	b
31	а	32	d	33	а	34	b	35	а
36	а	37	d	38	С	39	а	40	а
41	С	42	а	43	d	44	а	45	b
46	а	47	d	48	b	49	b	50	С
51	d	52	С	53	d	54	С	55	d
56	b	57	а	58	С	59	С	60	а
61	а	62	d	63	а	64	а	65	С
66	С	67	а	68	а	69	С	70	d
71	С	72	а	73	С	74	С	75	b
76	b	77	b	78	С	79	d	80	а
81	d	82	d	83	d	84	е	85	b
86	С	87	d	88	С	89	d	90	b
91	С	92	а	93	а	94	b	95	С
96	С	97	b	98	С	99	b	100	b
101	а	102	d	103	b	104	b	105	С
106	а	107	а	108	d				

## Halogen family

1	b	2	а	3	d	4	С	5	а
6	а	7	а	8	а	9	С	10	b
11	а	12	а	13	d	14	а	15	b
16	d	17	а	18	b	19	а	20	С
21	d	22	а	23	С	24	а	25	С
26	b	27	d	28	d	29	bd	30	d
31	d	32	а	33	d	34	а	35	d
36	а	37	b	38	а	39	d	40	а
41	d	42	b	43	d	44	а	45	a
46	а	47	b	48	а	49	d	50	b
51	а	52	а	53	d	54	а	55	а
56	d	57	а	58	С	59	а	60	d
61	b	62	С	63	а	64	а	65	d
66	b	67	b	68	b	69	С	70	С

71	b	72	а	73	а	74	b	75	d
76	b	77	b	78	С	79	С	80	b
81	b	82	С	83	d	84	d	85	d
86	а	87	а	88	а	89	d	90	b
91	d	92	С	93	С	94	а	95	а
96	С	97	b	98	b	99	С	100	а
101	а	102	С	103	d	104	b	105	d
106	е	107	b	108	а	109	а	110	а
111	С	112	С	113	а	114	а	115	a
116	С	117	b	118	b	119	b	120	b
121	d	122	а	123	b	124	а	125	a
126	С	127	d	128	а	129	а	130	С
131	d	132	а	133	а	134	а	135	a
136	С	137	d	138	b	139	d	140	а
141	d	142	d	143	d	144	b	145	а
146	а	147	а	148	b	149	С	150	a
151	d	152	b	153	d	154	а	155	С
156	b	157	b	158	а	159	С	160	d
161	d	162	а	163	b	164	а	165	а
166	С	167	С	168	С	169	b	170	a
171	а	172	b	173	С	174	С	175	b
176	b	177	С	178	b	179	а	180	d
181	а	182	а	183	а	184	С	185	а
186	а	187	С						

N	lob	le	ga	se	S

1	b	2	b	3	а	4	С	5	С
6	d	7	С	8	а	9	d	10	b
11	С	12	С	13	С	14	С	15	b
16	d	17	b	18	b	19	b	20	а
21	а	22	С	23	d	24	b	25	b
26	d	27	а	28	а	29	b	30	С
31	d	32	а	33	С	34	а	35	а
36	С	37	d	38	b	39	d	40	а
41	d	42	b	43	а	44	С	45	d
46	а	47	а	48	а	49	b	50	d
51	d	52	d	53	d	54	С	55	а
56	С	57	а	58	d	59	d	60	d

### **Critical Thinking Questions**

1	С	2	d	3	а	4	d	5	b
6	d	7	а	8	b	9	а	10	а
11	С	12	b	13	d	14	С	15	b

16	d	17	а	18	d	19	С	20	а
21	b	22	d	23	а	24	b	25	b
26	С	27	а	28	d	29	а	30	b
31	b	32	b	33	а	34	а	35	а
36	d	37	b	38	С	39	d	40	а
41	а	42	а	43	d	44	d	45	а
46	b	47	d	48	b	49	b	50	С
51	С	52	b	53	d	54	С	55	а
56	С	57	С	58	С	59	а	60	b
61	С	62	С	63	С	64	b	65	a
66	С	67	b	68	d	69	а	70	а
71	а								

Assertion and Reason										
1	b	2	b	3	а	4	а	5	а	
6	С	7	b	8	е	9	а	10	а	
11	е	12	а	13	а	14	а	15	а	
16	С	17	b	18	С	19	d	20	d	
21	а	22	b	23	d	24	d	25	b	
26	d	27	С	28	С	29	а	30	а	
31	а	32	С	33	b	34	а	35	а	
36	b	37	а	38	b	39	b	40	b	
41	а	42	С	43	а	44	d	45	С	
46	а	47	d	48	b	49	d	50	а	
51	е	52	b	53	d	54	b	55	С	
56	а	57	а	58	а	59	d	60	b	
61	а	62	b	63	С	64	b			

# Answers and Solutions

#### Alkali metals

1. (b) Element Na K  $IE_1$  496 419  $IE_2$  4562 3051

Sodium has higher I.E. because of smaller atomic size.

**2.** (c) Alkali metals are highly reactive metals. They react with

Alcohol – 
$$2C_2H_5OH + 2K \rightarrow 2C_2H_5OK + H_2$$

Water -  $2K + 2H_2O \rightarrow 2KOH + H_2$ 

Ammonia – 
$$K + (x + y)NH_3 \rightarrow \left[K(NH_3)_x\right]^+ + Ammoniated cation$$

 $[e(NH_3)_y]^-$ Ammoniated electron

But they do not react with kerosene.

- 4. (b) After removal of an electron the effective nuclear charge per electron increases hence the size decreases.
- **5.** (a) Alkali metals valence shell configuration =  $ns^1$
- **6.** (b) Element Li Na K Rb

Ionic radius 76 102 138 152 167 - (pm)

as the atomic no. increases the no. of shells increases hence, atomic radius increases.

(c) On moving down the group electropositive character increases. **8.** (a) Carnellite –  $KCl.\ MgCl_2.\ 6H_2O$ 

Cryolite -  $Na_3AlF_6$ 

Bauxite -  $(Al_2O_3.2H_2O)$ 

Dolomite - MgCO<sub>3</sub>. CaCO<sub>3</sub>

**10.** (d) Element – Li Na K Rb

Atomic radius (pm) 152 186 227 24 - 8

12. (b) Li is much softer than the other group I metals. Actually Li is harder then other alkali metals

13. (a) 
$$Cu^{+2} + 2e^{-} \rightarrow Cu$$
,  $E^{o} = +0.34 \text{ V}$   
 $Mg^{+2} + 2e^{-} \rightarrow Mg$ ,  $E^{o} = -2.37 \text{ V}$   
 $Na^{+} + e^{-} \rightarrow Na$ ,  $E^{o} = -2.71 \text{ V}$ 

- **14.** (d) Anhydrous form of  $Na_2CO_3$  does not decompose on heating even to redness. It is a amorphous powder called soda ash.
- 17. (c) Fehling's solution is a mixture of Alk.  $CuSO_4 + Na K$  tartarate (Rochelle salt)
- 19. (b)  $2K+2HCl \rightarrow 2KCl+H_2$  (violent reaction).
- **20.** (b) Although lattice energy of LiCl higher than NaCl but LiCl is covalent in nature and NaCl ionic there after, the melting point decreases as we move NaCl because the lattice energy decreases as a size of alkali metal atom increases (lattice energy  $\infty$  melting point of alkali metal halide)
- **22.** (b) It form calcium and magnesium complex with EDTA salt.
- 24. (a) LiOH < NaOH < KOH < RbOH Down the group basic character increases
- **25.** (d)  $Na_2CO_3$  .  $10H_2O \xrightarrow{\Delta} Na_2CO_3$  .  $H_2O \xrightarrow{\Delta}$  washing powder

 $Na_2CO_3 + H_2O \uparrow$ 

- **26.** (b)  $Na_2CO_3$ ,  $K_2CO_3$  and  $(NH_4)_2CO_3$  are soluble in water because hydration energy is more than lattice energy
- **29.** (c)  $K_2SO_4.Al_2(SO_4)_3.24H_2O$  potash alum it is a double salt.
- **31.** (d) It is a colourless gas.
- 32. (a)  $NaHCO_3 \rightarrow Na^+ + HCO_3^-$ (Salt of strong base & weak acid)  $OH^- + CO_2$
- **33.** (b)  $FeSO_4 . (NH_4)_2 SO_4 . 6H_2O$  Mohr's salt.
- **35.** (d)  $Ca^{+2} > Na^{+} > Mg^{+2} > Al^{+3}$
- **36.** (b)  $Li^+ + e^- \rightarrow Li$ ,  $E^o = -3.05 \ V$   $K^+ + e^- \rightarrow K$ ,  $E^o = -2.93 \ V$  $Ca^{+2} + 2e^- \rightarrow Ca$ ,  $E^o = -2.87 \ V$
- **37.** (a) Because their valence electrons are present in s- orbitals.
- **38.** (a)  $6Li + N_2 \rightarrow 2Li_3N$  Lithium nitride.

- **39.** (d) *Li*, *Na*, *K* are lighter than water but *Rb* is heavier than water.
- **42.** (c)  $KF + HF \rightarrow KHF_2 = K^+ + HF_2^-$
- **43.** (b) Cs > Rb > K > Na > Li Metallic character decreasing order.
- **45.** (d)  $2Rb + 2H_2O \rightarrow 2RbOH + H_2$ Li < Na < K < Rb < Cs

As we go down the group reactivity with  $H_2O$  increases.

- **48.** (b) Atomic number  $11 \rightarrow Na \rightarrow Na_2O$   $Na_2O + H_2O \rightarrow 2NaOH$ (base)
- **51.** (d) Generally ionic character decreasing from *LiCl* to *NaCl*.
- **52.** (c) In castner process *Na* metal is made of anode.
- 55. (a) Fajan's rule is applied.
- **57.** (a) Small atomic and ionic size leads to high electronegativity and hydration energy. Small atomic and ionic size leads to high electronegativity and hydration energy.
- **58.** (c) Mohr salt is  $(FeSO_4)(NH_4)_2SO_4.6H_2O$ .
- **60.** (a) Sodium thiosulphate is a reducing agent which convert metalic silver into silver salt.
- **64.** (a) In alkali metal group elements alkali means plant ash.
- **67.** (d)  $2Na + 2NH_3 \xrightarrow{\text{heat}} 2NaNH_2 + H_2$
- **68.** (a,b)  $2Na + \frac{1}{2}O_2 \xrightarrow{\text{moist air}} Na_2O$  $Na_2O + 2H_2O \xrightarrow{\text{moist air}} 2NaOH + H_2$ .
- **69.** (d)  $2KClO_3 \rightarrow 2KCl + 3O_2$ .
- **70.** (d) Due to free electron liquid ammonia becomes paramagnetic.
- **72.** (a) They possess highest atomic volume in their respective periods.
- **74.** (c)  $Fe(OH)_3$  is soluble in sodium hydroxide solution.
- **76.** (d) The cell involves the following reaction,

$$NaCl \Rightarrow Na^+ + Cl^-$$

At anode:  $2Cl^- \rightarrow 2Cl + 2e \rightarrow Cl_2$ 

At cathode :  $Na^+ + e \rightarrow Na$ 

 $Na + Hg \rightarrow \text{amalgam}$ 

At anode: Na – amalgam  $\rightarrow Na^+ + Hg + e$ 

At cathode :  $2H_2O + 2e \rightarrow H_2 \uparrow +2OH^-$ 

- **78.** (a) *Li* is a more reducing agent compare to other element.
- **79.** (b) Element *Li Na K Rb Cs*M.pt in *K* 4535 370.8 336.2 312
  301.5

- **80.** (a)  $2Na + 2HOH \rightarrow 2NaOH + H_2 \uparrow$  $2K + 2HOH \rightarrow 2KOH + H_2 \uparrow$
- **82.** (a) Alkali metal are good conductor of heat and electricity.
- **83.** (c) Potassium react with halogens (chlorine) to gives violet colour flame.
- **84.** (b) Mobility decreases from top to bottom because of the atomic size is increases.
- **85.** (c) Lithium shows digonal relationships with *Mg*.
- **86.** (c) K > Ca > C > Cl Electropositive character in decreasing order.
- **87.** (d)  $2NaCl \xrightarrow{\text{Electroly is}} 2Na + Cl_2 \xrightarrow{\text{Cathode}} Nach + Cl_2$
- **88.** (b) When sodium bicarbonate  $(NaHCO_3)$  is heated, sodium carbonate,  $CO_2$  and water are formed.

$$2NaHCO_3 \xrightarrow{\Delta} Na_2CO_3 + CO_2 \uparrow + H_2O$$
Sodium carbonate

- 89. (c) Alum is used for softning of water.
- **90.** (a) Only salts of (weak acid + strong base) and (strong acid + weak base) get hydrolysed (*i.e.*, show alkalinity or acidity in water).  $KClO_4$  is a salt of strong acid and strong base therefore it does not get hydrolysed in water.

$$KClO_4 \rightleftharpoons K^+ + ClO_4^-$$
;  $H_2O \rightleftharpoons OH^- + H^+$   
 $KOH \longrightarrow Strong \longrightarrow Strong$ 

- **91.** (c) Carbon dioxide does not help in burning, also it forms carbonate with alkali metals.
- **92.** (a) When carbonate are heated they decompose to form the oxide. Sodium carbonate and potassium carbonate do not decompose. The carbonate become more difficult to decompose as we go down the group.
- **93.** (c) Aluminium reacts with caustic soda to form sodium meta aluminate.

$$2Al + 2NaOH + 2H_2O \rightarrow 2NaAlO_2 + 3H_2 \uparrow$$
Sodium meta aluminate

- **94.** (a) Alkaline earth metals  $(ns^2)$  are denser than alkali metal  $(ns^1)$  because metallic bonding in alkaline earth metal is stronger.
- **95.** (c) Lithium is basic in nature and hence it is not amphoteric.
- **96.** (a) *CsOH* of the following is most basic in character due to increase electropositive character in a group of alkali.
- **97.** (a) Group I element are so highly electropositive that they emit electrons even when exposed to light (Photoelectric effect) and this character increase on moving down the group from lithium towards cesium.
- **98.** (b) Lithium form nitride on heating with nitrogen. Lithium nitride gives ammonia when

heated with  $H_2O$  . Ammonia gas form tetrammine copper complex with  ${\it CuSO}_4$  solution.

$$6Li + N_2 \rightarrow 2Li_3N$$

$$Li_3N + 3H_2O \rightarrow 3LiOH + NH_3$$

$$CuSO_4 + 4NH_3 \rightarrow [Cu(NH_3)_4]SO_4$$

**99.** (d) The given compound x must be  $CaCO_3$ . It can be explained by following reactions,

$$\begin{array}{c} CaCO_{3} \overset{\Delta}{\longrightarrow} CaO + CO_{2} \uparrow; \\ Ca(OH)_{2} + CO_{2} + H_{2}O \xrightarrow{} Ca(HCO_{3}) \\ \\ Ca(HCO_{3})_{2} \overset{\Delta}{\longrightarrow} CaCO_{3} + CO_{2} \uparrow + H_{2}O \end{array}$$

- 100. (c) According to Fajan's rule RbCl has greatest ionic character due to large ionic size of  $Rb^+$  ion.  $BeCl_2$  has least ionic (Maximum covalent) due to small size of  $Be^{+2}$  ion which has highly polarising.
- **105.** (c)  $2Na + 2NH_3 \rightarrow 2NaNH_2 + H_2$
- **110.** (d)  $2Na + 2H_2O \rightarrow 2NaOH + H_2$
- 112. (c) It reacts with alcohol to form sodium alkoxide  $2C_2H_5OH + 2Na \rightarrow 2C_2H_5ONa + H_2$
- 113. (a) Causticizing process (Gossage process) it is an old process and involves heating of 10% solution of  $Na_2CO_3$  with a little excess of milk of lime  $Ca(OH)_2$

$$Na_2CO_3 + Ca(OH)_2 \rightarrow CaCO_3 \downarrow +2NaOH$$

- **114.** (c)  $NaOH + CO \xrightarrow{150^{\circ}-200^{\circ}C} HCOONa$
- **115.** (a)  $2NaCl + 2H_2O \rightarrow 2NaOH + H_2 + Cl_2$
- **119.** (b) *NaOH* is a deliquescent white crystalline solid. It absorbs moisture from the atmosphere.
- **120.** (a)  $Na_2CO_3 + H_2O + 2SO_2 \rightarrow 2NaHSO_3 + CO_2$
- **123.** (c) NaOH + CaO is called soda lime 3:1
- 124. (c) Molten sodium is used as a coolant
- **126.** (c)  $AgBr + 2Na_2S_2O_3 \rightarrow Na_3[Ag(S_2O_3)_2] + NaBr$
- 128. (b)  $Na_2SO_3 + S \xrightarrow{NaOH} Na_2S_2O_3$
- **129.** (a,b)  $2NaCl + 2H_2O \rightarrow 2NaOH + Cl_2 + H_2$ Anode Cathod
- **132.** (b)  $Sn + 2NaOH + H_2O \rightarrow Na_2SnO_3 + 2H_2$
- **135.** (b)  $2NaCl + 2H_2O \xrightarrow{\text{Electroly is}} 2NaOH + Cl_2 + H_2$ Anode Cathod
- 136. (d)  $2NaCl \xrightarrow{\text{Electriccurrent}} 2Na^+ + 2Cl^-$ Cation Anion
- **138.** (a)  $HgCl_2 + 2NaOH \rightarrow HgO + 2NaCl + H_2O$

- 139. (a) Down's cell is used for the electrolysis of fused NaCl
- **142.** (c)  $Fe(OH)_3$  does not dissolve in NaOH
- **143.** (a) Castner's process used to obtain Na, by electrolysis of sodium hydroxide.
- **144.** (a) Excess of  $Na^+$  ion causes high B.P.
- **145.** (b) Ferric alum is  $(NH_4)_2SO_4.Fe_2(SO_4)_3.24H_2O$
- **146.** (d) When Na is heated in presence of air or  $CaO + H_2O \xrightarrow{} Ca(OH)$  burns to form sodium oxide and sodium personide.
- **148.** (d) Pyrolusite or Manganese dioxide  $(MnO_2)$  is a mineral of manganese.
- **149.** (c)  $CaCl_2$  bring down the melt temperature from 1075 K to 850 K

### Alkaline earth metals

- **2.** (d)  $CaSO_4 \cdot \frac{1}{2}H_2O$  or  $(CaSO_4)_2 \cdot H_2O$
- **3.** (a)  $CaCl_2$  because it is hygroscopic
- **7.** (d) Setting of plaster of paris is exothermic process

$$\begin{array}{c} \textit{CaSO}_{4}.\frac{1}{2}\textit{H}_{2}O \xrightarrow{\textit{H}_{2}O} \quad \textit{CaSO}_{4}.2\textit{H}_{2}O \\ & \xrightarrow{\text{Hardening}} \quad \textit{CaSO}_{4}.2\textit{H}_{2}O \\ & \xrightarrow{\text{Mono orthorhomb ic Gypsum}} \end{array}$$

The setting is due to formation of another hydrate

- 10. (a)  $MgCO_3 \xrightarrow{\text{Heat}} MgO + CO_2$ The metal whose oxide is stable, it's carbonate
- is unstable

  12. (d)  $MgCl_2 \xrightarrow{Electrolysis} Mg^{+2} + 2Cl$ (Molten)  $Cl_1 = 2Cl + 2l$ (Molten)  $Cl_2 = 2Cl + 2l$ (Molten)  $Cl_1 = 2Cl + 2l$

Anode – 
$$2Cl^- \rightarrow 2Cl + 2e^-$$
,  $Cl + Cl \rightarrow Cl_2$   
Cathode –  $Mg^{+2} + 2e^- \rightarrow Mg$ 

- **13.** (a) Because of small atomic size and high I.E. Be forms covalent chloride.
- 16. (d)  $BeSO_4$  is most soluble because hydration energy is more than lattice energy.  $BeSO_4 > MgSO_4 > CaSO_4 > SrSO_4 > BaSO$

Hydration energy decreases hence, solubility decreases.

- 19. (b)  $2(CaSO_4.2H_2O) \xrightarrow{120^{\circ}C} 2CaSO_4.H_2O + 3H_2O$ Gypsum 2 Plaster of paris
- **21.** (b) Lithopone  $(ZnS + BaSO_4)$  is used as a white pigment.
- 23. (d)  $Ca(OH)_2 + Cl_2 \rightarrow CaOCl_2 + H_2O$ Slaked lime

  Bleaching

  powder
- **24.** (a) Strontium  $\rightarrow$  Crimson or pink colour

- **26.** (b,c)  $Ca_3P_2 + 6H_2O \rightarrow 3Ca(OH)_2 + 2PH_3$  $K_3P + 3H_2O \rightarrow 3KOH + PH_3$
- 27. (d)  $CaCl_2 \rightarrow Ca^{+2} + 2Cl^-$ Cathode:  $Ca^{+2} + 2e^- \rightarrow Ca$ Anode:  $2Cl^- \rightarrow 2e^- + Cl_2$
- 28. (d) Element Mg Al Si

  Atomic radii 1.60 1.43 1.32 1.28
  (Å)-

as we move across the period nuclear charge increases, hence, size decreases.
 30. (b) MgCl<sub>2</sub>.6H<sub>2</sub>+5MgO+xH<sub>2</sub>O →

 $MgCl_2.5MgO.xH_2O$ 

- Magnesia cement or sorrel cement 31. (d)  $ZnS + BaSO_4$  is lithopone used as white
- pigment. **36.** (d) Aqueous  $CaCl_2$  or hydrated  $CaCl_2$  can not act as dehydrating agent.
- **38.** (d) As we go down the group electropositive character increases because I.E. decreases. Ba is most electropositive element in the group.
- **39.** (d) Due to the inert pair effect.
- **40.** (a) Element Be Al Electronegativity 1.5 1.5
- 41. (c) Be > Mg > Ca > Sr > BaOn moving down the group lattice energy remains almost constant as the sulphate is so big that small increase in the size of the cations from Be to Ba does not make any difference. However the hydration energy decreases from  $Be^{+2}$  to  $Ba^{+2}$ . This causes decrease in the solubility of the sulphates as
- **42.** (a) Element Be Mg Ca Sr Ba

  Electrode potential 1.70 2.37 2.87 2.89

-2.90 **43.** (a) Element - *Mg Ca Sr Ba* I.E - 737 590 549 503

the ionic size increases.

- 44. (a) Be due to diagonal relationship
- **45.** (a)  $K^+$  is highly soluble because of high hydration energy.
- 47. (b) MgO  $Al_2O_3$   $SiO_2$   $P_2O_5$ Basic Amphoteric Acidic  $MgO + H_2O \rightarrow Mg(OH)_2$  Base or alkali
- **48.** (d) Duralium (Al = 95%, Cu = 4%, Mn = 0.5%, Mg = 0.5%)

being light, tough and durable is used for the manufacture of aeroplanes and automobile parts.

- **49.** (c) Na K Ba Ca Sr Yellow Pale Apple Brick Crimson violet green red
- **51.** (a) Magnesium burns with an intense light. Therefore *Mg* is used in flash bulbs for photography, fireworks and signal fires.
- **53.** (b)  $CaO + CO_2 \rightarrow CaCO_3$  $CaO + H_2O \rightarrow Ca(OH)_2$
- **54.** (a) When water is added to cement an exothermic reaction occurs. During this process, the cement reacts with water to form a gelatinous mass which slowly sets into a hard mass having three dimensional network structure involving -Si O Si and -Si O Al chains.
- **55.** (b) CaO (quick lime)  $Ca(OH)_2$  (slaked lime)  $Ca(OH)_2 + H_2O$  an aqueous suspension of  $Ca(OH)_2$  in water is called lime water.  $CaCO_3$  (lime stone).
- 57. (d) Lime stone  $CaCO_3$ Clay – silica and alumina Gypsum –  $CaSO_4.2H_2O$
- **59.** (b) Because hydration energy decreases down the group.
- **63.** (d) Be does not react with water.
- (ii) High electronegativity
  (iii) Absence of *d* orbitals
- **65.** (a)  $\underbrace{Ba(OH)_2 > Sr(OH)_2 > Ca(OH)_2 > Mg(OH)_2}_{\text{Solubility decreasing order.}} > Mg(OH)_2$
- **66.** (d) Solubility increasing top to bottom.
- **67.** (a) Be to Ba ionic character increasing.
- **70.** (a)  $Ca + 2H_2O \rightarrow Ca(OH)_2 + H_2$   $CaH_2 + 2H_2O \rightarrow Ca(OH)_2 + 2H_2$
- **72.** (a) They are denser than alkali metals because they can be packed more tightly to their greater charge and smaller radii.
- **76.** (d)  $Be(OH)_2 < Mg(OH)_2 < Ca(OH)_2 < Sr(OH)_2 < Ba(OH)_2$ On moving down the group basic character increases.
- 77. (b)  $Mg(OH)_2$  Mg is most electropositive element amongst the given elements.
- **78.** (d) Lime stone =  $CaCO_3$ Quick lime = CaOSlaked lime =  $Ca(OH)_2$

- **79.** (c) As we go down the group I.E. decreases. Hence, *Ba* can easily give electrons. Therefore strongest reducing agent.
- **80.** (c)  $CaSO_4$ .  $\frac{1}{2}H_2O + \frac{1}{2}H_2O \to CaSO_4$ .  $2H_2O$ Plaster of paris
- **82.** (a)  $BeCl_2 < MgCl_2 < CaCl_2 < BaCl_2$ As we go down the group I.E. decreases. Hence ionic character increases.
- **83.** (c)  $MgCl_2.6H_2O \xrightarrow{\text{heat}} MgO + 5H_2O + 2HCl$
- **84.** (a) The solubility of hydroxides of alkaline earth metals in water increases on moving down the group.
- **85.** (c) Bleaching action of  $Cl_2$  in moist condition is permanent.  $Cl_2 + H_2O \rightarrow HCl + HClO$

$$\frac{HClO \to HCl + O}{Cl_2 + H_2O \to 2HCl + O}$$

- **86.** (d) The solubility of hydroxides of alkaline earth metals in water increases on moving down the group.
- **87.** (c)  $CO_2$  escapes simultaneously.
- **88.** (d) It consist of high lattice energy and lowest hydration energy.
- **90.** (a) It is a *s*-block elements.
- **91.** (b) Barium  $Ba_{56}$  is a alkali earth metal.
- **92.** (c) Thermal stability increasing from top to bottom.
- **94.** (b) On moving down the group; Lattice energy decreases with increase in size of cation.
- **96.** (d)  $BaSO_4$  is sparingly soluble in water because the solubility of second group sulphates decreases with increasing atomic size. Because of hydration energy decreases.
- **97.** (d) Berylium because of small atomic size and high ionization energy.
- **99.** (a) Mg < Ca < Sr < Ba < RaIonic nature increases

as we go down the group ionic nature increases because I.E. decreases.

- 100. (b)  $CaSO_4 . 2H_2O$  Gypsum  $CaSO_4 . \frac{1}{2} H_2O$  Plaster of paris
- **101.** (b)  $Ca + \frac{1}{2}O_2 \to CaO$ .
- **102.** (c) Calcium cyanamide is the slow acting nitrogenous fertilizer as it decompose very slowly.

$$CaNCN + 2H_2O \rightarrow CaCO_3 + NH_2CONH_2$$

$$NH_2CONH_2 + H_2O \rightarrow CO_2 + 2NH_3$$
  
 $NH_3 \xrightarrow{\text{Nitrifying}} \text{Soluble nitrates} \rightarrow \text{Plants}$ 

103. (a) Plaster of paris  $[(CaSO_4)_2H_2O]$  is used in surgery for setting of bones, dentistry and manufacturing of statues. It is prepared as follows.

$$2 CaSO_4.2 H_2 O \xrightarrow{125\,{}^{\circ}C} \rightarrow (CaSO_4)_2. H_2 O + 3 H_2 O$$
Plaster of paris

- **104.** (b) Due to electropositive and reactive in nature, magnesium is readily converted into positive ions on contact with iron pipes and hence, iron pipes remains as it is.
- 105. (c) A binary compound is one made of two different elements. These can be one of each element such as CuCl or FeO. These can also be several of each element such as  $Fe_2O_3$  or  $SnBr_4$ . Metal which have variable oxidation number can form more than one type of binary compound like Fe shows the oxidation state +2 and +3. Hence it forms two type of binary compound e.g.,  $FeCl_2$ ,  $FeCl_3$ .
- 106. (a) Diagonal relationship: elements of  $2^{nd}$  period often show resemblance to the element of the IIIrd period diagonaly placed to it. This type of behaviour is called as diagonal relationship Li shows the diagonal relationship with Mg.

**107.** (a) 
$$MgCl_2 + 2NaHCO_3 \rightarrow Mg(HCO_3)_2(aq) + 2NaCl$$
 
$$Mg(HCO_3)_2(aq) \xrightarrow{\Delta} MgCO_3 \downarrow + H_2O + CO_2 \uparrow$$
 (White ppt.)

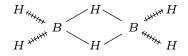
**108.** (c) We know that  $MgCl_2.6H_2O \xrightarrow{Heat} MgCl_2 + 6H_2O$ 

Thus in this reaction magnesium dichloride is produced.

- **109.** (d) Magnesium burns in *CO* to produce  $Mg + CO \rightarrow MgO + C$
- **110.** (d) Sorel's cement is,  $MgCl_2.5MgO.xH_2O$
- 111. (b) Colemnite is a mineral of boron having composition as  $Ca_2B_6O_{11}.5H_2O$ .

### **Boron family**

- 3. (a)  $B_2O_3 + 3C + 3Cl_2 \rightarrow 2BCl_3 + 3CO$  $BCl_3$  is obtained by passing chlorine over the heated mixture of  $B_2O_3$  and powdered charcoal.
- **6.** (d)  $B_2H_6$  has two types of B-H bonds



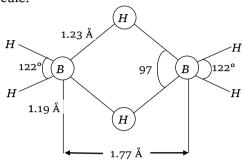
119 pm

134 pm

#### B H (Terminal bond)

B H (Bridge bond)

12. (b) Dilthey in 1921 proposed a bridge structure for diborane. Four hydrogen atoms, two on the left and two on the right, known as terminal hydrogens and two boron atoms lie in the same plane. Two hydrogen atoms forming bridges, one above and other below, lie in a plane perpendicular to the rest of molecule.



- **15.** (c)  $2H_3BO_3 \rightarrow B_2O_3 + 3H_2O$ .
- **16.** (a,c,d)  $Al_2Cl_6$ ,  $In_2Cl_6$ ,  $Ga_2Cl_6$
- **17.** (a) Liquified *Ga* expand on solidification *Ga* is less electropositive in nature, It has the weak metallic bond so it expand on solidification.
- **18.** (d)  $Al_2Cl_6 + 12H_2O = 2[Al(H_2O)_6]^{3+} + 6Cl^{-1}$
- **19.** (e)  $B_4C$  is the hardest substance along with diamond.
- **20.** (a) Borazine  $B_3N_3H_6$ , is isoelectronic to benzene and hence, is called inorganic benzene some physical properties of benzene and borazine are also H nilar.

- **21.** (c) Except  $B(OH)_3$  all other hydroxide are of metallic hydroxide having the basic nature  $B(OH)_3$  are the hydroxide of nonmetal showing the acidic nature.
- **22.** (c) Moissan boron is amorphous boron, obtained by reduction of  $B_2O_3$  with Na or Mg. It has 95-98% boron and is black in colour.
- **23.** (d) Boron form different hydride of general formula  $B_nH_{n+4}$  and  $B_nH_{n+6}$  but  $BH_3$  is unknown.

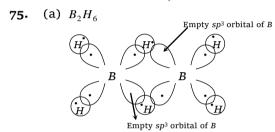
- **24.** (c) Alumina is amphoteric oxide, which reacts acid as well as base.
- **25.** (a) Al is the most abundant metal in the earth crust.
- **29.** (a)  $AlCl_3.6H_2O \xrightarrow{\Delta} Al(OH)_3 + 3HCl + 3H_2O$ Thus  $AlCl_3$  can not be obtained by this method
- **30.** (d) Amphoteric substance can react with both acid and base.
- **33.** (c)  $2Al + 6HCl \rightarrow 2AlCl_3 + 3H_2$
- **34.** (c)  $Al \rightarrow III \text{ group} \rightarrow Forms <math>Al_2O_3$
- **35.** (d)  $2KOH + 2Al + 2H_2O \rightarrow 2KAlO_2 + 3H_2$
- 37. (c)  $Na_2CO_3 + H_2O \rightarrow 2NaOH + CO_2$  $2NaOH + 2Al + 6H_2O \rightarrow 2Na[Al(OH)_4] + 3H_2$
- **41.** (c)  $B(OH)_3 \Rightarrow H_3BO_3$  Boric acid  $Al(OH)_3 \Rightarrow$  Amphoteric
- **45.** (b)  $FeSO_4.(NH_4)_2SO_4.6H_2O$
- **46.** (b)  $Al_2O_3$  is an amphoteric oxide.
- **47.** (c) Aluminium oxide is highly stable therefore, it is not Reduced by chemical reactions.
- **48.** (d) Aluminium is used as reducing agent in metallurgy.
- **49.** (a) Al is used as reducing agent in thermite process.
- **50.** (c) In Goldschmidt aluminothermic process, thermite contains 3 parts of  $Fe_2O_3$  and 1 part of Al.
- 51. (c) For the purification of red bauxite which contains iron oxide as impurity → Baeyer's process. For the purification of white bauxite which contains silica as the main impurity Serpeck's process.
- 52. (b) In Hall's process  $Al_2O_3.2H_2O + Na_2CO_3 \rightarrow 2NaAlO_2 + CO_2 + 2H_2O$   $2NaAlO_2 + 3H_2O + CO_2 \xrightarrow{333 \ K} \rightarrow$   $2Al(OH)_3 \downarrow + Na_2CO_3$   $2Al(OH)_3 \xrightarrow{1473 \ K} \rightarrow Al_2O_3 + 3H_2O$
- **54.** (d) Cryolite  $Na_3AlF_6$ 
  - (1) Decreases the melting point of alumina(2) Increases conductivity of the solution
- **55.** (b) Cryolite  $Na_3AlF_6$  is added
  - (1) To decrease the melting temp from 2323 K to 1140 K
  - (2) To increase the electrical conductivity of solution
- **61.** (d) Iron oxide impurity Baeyer's process Silica impurity Serpeck's process
- **64.** (b) Cryolite is added to lower the melting point of alumina and to increase the electrical conductivity.
- **65.** (c) The purification of alumina can be done by Baeyer's process.

- **67.** (c) In electrolytic method of obtaining aluminium from purified bauxite, cryolite is added to charge because it reduces the melting point of Bauxite (from  $1200\,^{o}C$  to  $800\,^{o}-900\,^{o}C$ ) and also it increases electrical conductivity of mixture.
- **68.** (a) Hoop's process  $\Rightarrow$  Purification of Al Hall and Heroult process  $\Rightarrow$  Reduction of  $Al_2O_3$

Baeyer's and Serpeck's process  $\Rightarrow$  Concentration of Bauxite ore

74. (a) 
$$H \searrow H \searrow H$$

3c-2e:B-H-B; 2c-2e:H-B-H



**76.** (b) Pure alumina is a bad conductor of electricity and the fusion temperature of pure alumina is about 2000°C and at this temperature when the electrolysis is carried of fused mass the metal formed vapoureses as the boiling point of *Al* is 1800°C.

To overcome this difficulty,  $Na_3AlF_6$  and  $CaF_2$  are mixed with alumina.

77. (a) Concentration of Lewis acid of boron tri halides is increased in following order.  $BF_3 < BCl_3 < BBr_3 < Bl_3$ .

### Carbon family

- 3. (d) It react with alkali as well as acid.
- **6.** (a) Among alkali metal carbonates only  $Li_2CO_3$  decomposes.  $Li_2CO_3 \xrightarrow{\Delta} Li_2O + CO_2 \uparrow$
- 7. (b) Propyne can be prepared by the hydrolysis of magnesium carbide.  $Mg_2C_3 + 4H_2O \rightarrow CH_3C \equiv CH + 2Mg(OH)_2$
- 10. (d) Generally red lead decompose into PbO and  $O_2$ .
- 11. (c)  $CO_2$  is acidic oxide and thus more effectively absorbed by an alkali.
- **12.** (b)  $CaC_2$  have one sigma and two  $\pi$  bond.
- **13.** (d) *C* and *Si* are non-metal and *Pb* is a metal.
- **16.** (a)  $SiO_2 + 2Mg \rightarrow Si + 2MgO$ .

- 17. (b) Generally IV group element shows catenation tendency and carbon has more catenation power.
- **18.** (b) Metal oxides or some salts are fused with glass to imported colour of glass.
- **19.** (d)  $Al_2(CO_3)_3$  is less soluble in water than  $Na_2CO_3$   $ZnCO_3$ .
- **20.** (d) The inert pair effect is most prominent in *Pb* because from top to bottom due to increase in number of shells.
- 25. (c)  $Co + NaOH \xrightarrow{200^{\circ}C} HCOONa$ Sod. formate
- 27. (c) Sodium oxalate react with conc.  $H_2SO_4$  to form CO and  $CO_2$  gas.
- **33.** (d) It is hydrolysed with water to form a  $Si(OH)_4$ .
- **35.** (b) When hydrogen peroxide react with *PbS* then they form  $PbSO_4$ .
- 36. (b) Grey tin is very brittle and easily crumbles down to a powder in very cold climates.
  Cross tin A White tip

Grey tin 

⇒ White tin (Tetragonal)

The change of white tin to grey tin is accompanied by increase in volume. This is called tin disease or tin plague.

- 37. (c) Solid  $CO_2$  is knows as dry ice because it evaporates at  $-78^{\circ}C$  without changing in the liquid state.
- **38.** (b) Zeolite have  $SiO_4$  and  $AlO_4$  tetrahedrons linked together in a three dimensional open structure in which four or six membered ring predominate. Due to open chain structure they have cavities and can take up water and other small molecules.
- **39.** (b) Crook's glass is a special type of glass containing cerium oxide. It does not allow the passage of ultra violet ray and is used for making lenses.
- **40.** (b) Inert pair effect become significant for the  $6^{th}$  and  $7^{th}$  period of *p*-block element.
- **41.** (a) Carbon suboxide has linear structure with C-C bond length equal to 130  $\mathring{A}$  and C-O bond length equal to 120  $\mathring{A}$ .

$$O = C = C = C = O \Leftrightarrow O^{-} - C \equiv C - C \equiv O^{+}$$

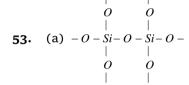
- **42.** (c)  $Pb_3O_4$  is a mixed oxide. It can be represented as  $2PbO PbO_2$ .
- **43.** (b) Noble gases are found in very minute amount in atmosphere. These are separated from each other by using coconut charcoal. Which adsorb different gas at different temperature.
- **44.** (c) Lapis Lazuli is a rock composed mainly of the following mineral, lazurite, hauynite sodalite, nosean, calcite, pyrite, lapis lazuli is actually sulphur containing, sodium aluminium silicate having chemical composition  $3Na_2O.3Al_2.6SiO_2.2Na_2S$ .

- **45.** (d) In carbon family stability +2 oxidation state increases on moving down the group in the periodic table with an increase in atomic number due to screening effect.
- **46.** (c) Tin is oxidised to meta stannic acid when it is treated with nitric acid.

 $Sn + 4HNO_3 \rightarrow H_2SnO_3 + 4NO_2 + H_2O$ 

- **47.** (c) Pb + Sn
- **49.** (d) Three dimensional sheet structures are formed when three oxygen atoms of each  $[SiO_4]^{4-}$  tetrahedral are shared.
- **50.** (a)  $Pb_3O_4 \Rightarrow \text{Red lead (Sindhur)}$
- **51.** (c) White lead  $\Rightarrow 2PbCO_3.Pb(OH)_2$
- **52.** (c) Organic acids dissolve lead in presence of oxygen

$$Pb + 2CH_3COOH + \frac{1}{2}O_2 \rightarrow Pb(CH_3COO)_2 + H_2O$$



- **55.** (a)  $S^2P^2$  Total 4 valence electrons  $\Rightarrow$  IV group
- **56.** (c)  $PbCl_2$  is most ionic because on going down the group the metallic character increases and also the inert pair effect predominates.
- **58.** (b) Type metal Pb = 82%, Sb = 15%, Sn = 3%
- **60.** (b) Sugar of lead  $(CH_3COO)_2Pb \Rightarrow$  lead acetate
- **63.** (d)  $Pb \Rightarrow 11.34 \ q/ml$  Heaviest
- **64.** (c)  $Pb_3O_4$  is a mixed oxide of  $2PbO + PbO_2$
- **67.** (c) Boron (B), Si, Ge, As, Sb, and At are the metalloid elements. Bismuth (Bi) and tin (Sn) are metals while carbon (C) is non-metal.
- **68.** (a)  $Al_4C_3 + 12H_2O \rightarrow 3CH_4 + 4Al(OH)_3$
- **69.** (b) Glass being a mixture of sodium and calcium silicates reacts with hydrofluoric acid forming sodium and calcium fluorosilicates respectively.

 $Na_2SiO_3 + 3H_2F_2 \rightarrow Na_2SiF_4 + 3H_2O$  $CaSiO_3 + 3H_2F_2 \rightarrow CaSiF_4 + 3H_2O$ 

The etching of glass is based on these reactions.

### Nitrogen family

- 3. (b)  $FeSO_4 + NO \rightarrow FeSO_4.NO$
- **4.** (b)  $HPO_3$ , metaphosphoric acid



**6.** (a) White phosphorus is soluble in  $CS_2$  whereas red phosphorus is insoluble in it.

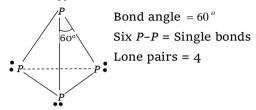
7. (d)  $H_4P_2O_7$  pyrophosphoric acid

$$HO - P - O - P - OH$$

$$OH OH$$

Tetrabasic (4-OH groups)

- **8.** (b)  $P_4 + 3NaOH + 3H_2O \rightarrow PH_3 + 3NaH_2PO_2$  (White) Phosphine Sod. hypophosph ite
- **9.** (a) *NCl*<sub>5</sub> is not known because of absence of *d*-orbitals in nitrogen.
- 11. (a,d)  $P_4$  molecule



- 12. (b)  $NH_4NO_3 \xrightarrow{\Delta} 2H_2O \uparrow + N_2O \uparrow$ Nitrous oxide (Laughing gas)
- 13. (a) Birkeland Eyde process

  Dinitrogen is prepared commercially from air by liquification and fractional distillation.

  When liquid air is allowed to distil, dinitrogen having lower b.pt (77K) distils over first leaving behind liquid oxygen (bpt 90K). World wide production of dinitrogen from liquid air is more than 50 million tonns per year.
- 14. (b)  $NH_4 NO_3 \xrightarrow{\Delta} 2H_2O \uparrow + N_2O \uparrow$   $NaNO_3 \xrightarrow{\Delta} NaNO_2 + O_2 \uparrow$   $2AgNO_3(s) \rightarrow 2Ag(s) + 2NO_2(g) + O_2(g)$ Lunar caustic  $2Pb(NO_3)_2 \rightarrow 2PbO + 4NO_2 \uparrow + O_2 \uparrow$
- **16.** (b)  $P_4 + 3NaOH + 3H_2O \rightarrow PH_3 + 3NaH_2PO_2$ White Phosphine
- 17. (d) NP As Sb Bi Non-metals Metalloids Meta
- 18. (b) O | P | P | HO OH OH
  - 3 *OH* groups are present hence it is tribasic.
- 19. (c) Nitrous acid behaves as reducing as well as an oxidising agent. It reduces potassium permanganate, potassium dichromate,  $H_2O_2$  and other strong oxidising agents. It oxidises strong reducing agents such as hydroiodic acid, sulphurous acid etc.

It oxidises  $Fe^{+2}$  into  $Fe^{+3}$  in acidic medium;  $Fe^{+2} + HNO_2 + H^+ \rightarrow Fe^{+3} + NO + H_2O$ 

It reduces acidified  $KMnO_4$ .

$$2KMnO_4 + 3H_2SO_4 + 5HNO_2 \rightarrow$$

$$K_2SO_4 + 2MnSO_4 + 3H_2O + 5HNO_3$$

**20.** (a) 
$$2KNO_3 \xrightarrow{\Delta} 2KNO_2 + O_2$$

23. (b) 
$$NH_4Cl + KNO_2 \rightarrow NH_4NO_2 + KCl$$

$$\downarrow N_2 + 2H_2O$$

**24.** (d) 
$$2HNO_3 \rightarrow N_2O_5 + H_2O_5$$

**25.** (c) 
$$2Ca_3(PO_4)_2 + 6SiO_2 \xrightarrow{1770\text{K}} 6CaSiO_3 + P_4O_{10}$$
  
 $P_4O_{10} + 10C \xrightarrow{1770\text{K}} P_4 + 10CO$ 

**26.** (c) 
$$P_4 + 5O_2 \rightarrow 2P_2O_5$$
 
$$P_2O_5 + 3H_2O \rightarrow 2H_3PO_4$$

**27.** (c) 
$$H_3PO_4 = H^+ + H_2PO_4^-$$
  
 $H_2PO_4^- = H^+ + HPO_4^{2-}$   
 $HPO_4^{2-} = H^+ + PO_4^{3-}$ 

**30.** (c) 
$$(NH_4)_2 Cr_2 O_7 \xrightarrow{\Delta} Cr_2 O_3 + N_2 + 4H_2 O$$

**31.** (d) 
$$4HNO_3 \rightarrow 4NO_2 + O_2 + 2H_2O_3$$

**32.** (c) Because of its very low ignition temperature (303K) it is always kept under water.

33. (d) 
$$NH_3$$
 when dissolved in water forms 
$$NH_3 + H_2O \rightarrow NH_4^+ + OH^- \implies NH_4OH$$

34. (a,b) Stability of + 3 oxidation states increases on account of inert pair effect.Reducing character of hydrides increases down the group because bond dissociation energy decreases down the group.

35. (a) Haber's process 
$$\rightarrow$$
 Industrial process
$$N_2 + 3H_2 \xrightarrow{Fe - Mo} 2NH_3$$

$$\xrightarrow{650 - 800 \text{ K}} 200 - 350 \text{ atm}$$

**36.** (a)  $P_4 + 5O_2 \rightarrow P_4O_{10}$ ; white phosphorus gets easily oxidized because it is highly reactive.

**38.** (d)  $N_2O$  is itself non-combustible but supports combustion.  $S + 2N_2O \rightarrow SO_2 + 2N_2$ 

**39.** (b) When  $N_2O$  is inhaled in moderate quantities, it produces hysterical laughter, hence the name laughing gas.

**40.** (c) 
$$2NO_2 + H_2O$$

$$\Rightarrow HNO_2 + HNO_3$$

When dissolved in water, gives a mixture of nitrous acid and nitric acid.

$$N_2O_4 + H_2O \rightarrow HNO_2 + HNO_3$$

 $HNO_3 + HNO_2 + 2NaOH \rightarrow NaNO_2 + NaNO_3 + 2H_2O$ 

**41.** (b) 
$$4Zn + 10HNO_3 \rightarrow 4Zn(NO_3)_2 + 5H_2O + N_2O_3$$

**42.** (a)  $(HPO_3)_n$  Polymetaphosphoric acid

**43.** (b) Superphosphate of lime – It is a mixture of calcium dihydrogen phosphate and gypsum and is obtained by treating phosphatic rock will conc.  $H_2SO_4$ 

$$\begin{array}{l} Ca_3(PO_4)_2 + 2H_2SO_4 + 5H_2O \rightarrow \\ Ca(H_2PO_4)_2.2H_2O + 2CaSO_4.2H_2O \\ \text{Superphosp hate of lime} \end{array}$$

**44.** (d) 
$$3NaOH + H_3PO_4 \rightarrow Na_3PO_4 + 3H_2O_4$$

**46.** (d) 
$$4NH_3 + 5O_2 \xrightarrow{Pt} 4NO + 6H_2O$$

NO is used in the preparation of  $HNO_3$  $2NO + O_2 \rightarrow 2NO_2$ ;  $4NO_2 + 2H_2O + O_2 \rightarrow 4HNO_3$ 

**47.** (d) 
$$P_4 + 20HNO_3 \rightarrow 4H_3PO_4 + 20NO_2 + 4H_2O$$

**49.** (b) 
$$P_2O_3$$
  $A_2O_3$   $B_2O_3$   $Bi_2O_3$  Alkalin

**50.** (b) 
$$P_2O_3 + 3H_2O \rightarrow 2H_3PO_3$$

(a) 
$$NF_3$$
 $F$ 
 $F$ 

51.

It is least basic because of the high electronegativity of 3F atoms. The lone pair present on nitrogen atom is not easily available for donation.

**52.** (d) 
$$3H_2O + PCl_3 \rightarrow H_3PO_3 + 3HCl$$

53. (c) Due to less reactivity of red phosphorus

**54.** (d)  $NO_2$  brown coloured gas.

**55.** (d)  $N_2O_5$  is an anhydride of  $HNO_3$  $2HNO_3 \rightarrow N_2O_5 + H_2O$ 

Therefore, it can act only as oxidising agent.

**56.** (c) 
$$NH_4NO_2$$
  $\rightleftharpoons$   $NH_4^+$  +  $NO_2^ x+4=+1$   $x-4=-1$   $x=1-4=-3$   $x=+3$ 

57. (b) 
$$P_4 O_8$$
  
 $4x + (-2 \times 8) = 0$   
 $4x - 16 = 0$   
 $x = \frac{16}{4} = +4$ 

**58.** (c) 
$$NH_2OH$$
  
 $x + 2 + (-2) + 1 = 0$   
 $x + 2 - 2 + 1 = 0$   
 $x = -1$ 

**60.** (c) 
$$NH_3 > PH_3 > AsH_3 > SbH_3$$
  
On moving down the group atomic size increases and availability of lone pair decreases. Hence, basic character decreases.

**61.** (a) 
$$PH_3 > AsH_3 > SbH_3 > BiH_3$$
  
On moving down the group bond energy decreases. Hence, stability decreases.

**62.** (d) Due to absence of *d*-orbitals in 
$$N$$
 atom, it cannot accept electrons from  $H_2O$  for hydrolysis of  $NF_3$ .

**63.** (b) 
$$NH_3$$
 is most thermally stable hydride. Hence, electrolysis temperature is maximum.

**64.** (a) Phosphorus is kept in water due to it burt at 
$$30^{\circ} C$$
.

**66.** (c) 
$$BiCl_3 + H_2O \rightarrow BiOCl + 2HCl$$

**67.** (c) When the black ppt. of 
$$Bi_2S_3$$
 is dissolved in 50%  $HNO_3$  and a solution of  $NH_4OH$  is added. A white ppt. of  $Bi(OH)_3$  is obtained.

70. (b) Bismuth does not show allotropy other elements show allotropy. Nitrogen 
$$\rightarrow \alpha$$
-nitrogen and  $\beta$ -nitrogen (solid crystalline forms) Phosphorus  $\rightarrow$  White, Red and Black forms Arsenic  $\rightarrow$  Yellow and Grey forms Antimony  $\rightarrow$  Yellow and Grey forms

**72.** (a) 
$$NH_3$$
 is a strongest base because Lone pair is easily available for donation.

**74.** (b) Hydride 
$$NH_3$$
  $PH_3$   $AsH_3$   $SbH_3$   $BiH_3$ 

Boling point 238.5 185.5 210.6 254.6

**75.** (a) 
$$NCl_3$$
 is highly reactive and unstable. Hence it is explosive.

**76.** (b) 
$$N_2O_3$$
  $P_2O_3$   $As_2O_3$   $Sb_2O_3$   $Bi_2O_3$ 
Acidic Oxides Amphoteri Basic

Acidic character decreases down the group

77. (c) 
$$SbCl_2$$
 is not exists because  $V^{th}$  group elements normally show +3 and +5 oxidation state.

**78.** (b) 
$$NH_4Cl + NaNO_2 \rightarrow NH_4NO_2 + NaCl$$

$$NH_4NO_2 \xrightarrow{\text{heat}} N_2 + 2H_2O$$
(g) (l)

**79.** (c) 
$$NH_4NO_2 \rightarrow N_2 + 2H_2O$$

**80.** (d) 
$$6Li + N_2 \rightarrow 2Li_3N$$
 Lithium nitride  $3Mg + N_2 \rightarrow Mg_3N_2$  Magnesium nitride

**81.** (d) 
$$N = N$$
 bond energy is very high  $945 \, kJ \, mol^{-1}$ .

**83.** (d) 
$$N_7 \rightarrow 1s^2, 2s^2, 2p^3$$
 d-orbitals are absent in nitrogen.

**85.** (d) 
$$NH_4NO_3 \xrightarrow{\text{heat}} N_2O + 2H_2O$$
 (Laughing gas)

**86.** (d) 
$$NH_2OH + HNO_2 \rightarrow H_2 N_2 O_2 + H_2O_1$$

**87.** (c) 
$$N_2O$$
 is a linear molecule

**88.** (b) 
$$2HNO_2 \rightarrow H_2O + N_2O_3$$

**89.** (d) 
$$2HNO_3 \rightarrow H_2O + N_2O_5$$

**90.** (c) 
$$2Pb(NO_3)_2 \rightarrow 2PbO + 4NO_2 + O_2$$

**91.** (a) 
$$2Pb(NO_3)_2 \rightarrow 2PbO + 4NO_2 + O_2$$

**92.** (b) 
$$3Cu + 8HNO_3 \rightarrow 3Cu(NO_3)_2 + 4H_2O + 2NO_3$$

**93.** (d) In upper atmosphere *NO* is formed by lightning flash.

$$N_2 + O_2 = 2NO$$

**96.** (c) 
$$2NO + O_2 \rightarrow 2NO_2$$

**100.** (d) 
$$2NO_2 + H_2O \rightarrow HNO_3 + HNO_2$$

**101.** (d) 
$$C_{12}H_{22}O_{11} \xrightarrow{\text{conc.} HNO_3} \xrightarrow{COOH} + H_2O$$

**102.** (b) 
$$4NH_3 + 5O_2 \xrightarrow{Pt} 4NO + 6H_2O$$

103. (d)  $HNO_2$  can be either reduced to nitric oxide (NO) or oxidised to nitric acid and hence it acts both as an oxidising as well as a reducing agent.

$$2HNO_2 \rightarrow 2NO + H_2O + [O]$$
  
 $HNO_2 + [O] \rightarrow HNO_3$ 

106. (d)  $NH_3$  is highly volatile compound. When vapourized, liquid ammonia causes intense cooling. Hence used as a coolant in ice factories and cold storages.

**107.** (d) 
$$N_3 H \Rightarrow N_3^- + H^+$$

**108.** (a) *d*-orbitals are absent in nitrogen.

109. (d) Phosphide ion Chloride ion

$$(P^{3-})$$
  $(Cl^{-})$  as 18 18

Total electrons 18 18 
$$P^{3-}$$
 and  $Cl^{-}$  are isoelectronic.

110. (a) Due to the less reactivity.

**116.** (a) 
$$P_4 + 3H_2O + 3NaOH \rightarrow PH_3 + 3NaH_2PO_2$$
  
Phosphine Sod. hypophosph ite

Oxidation

$$\stackrel{0}{P_4} + 3H_2O + 3NaOH \rightarrow \stackrel{-3}{PH_3} + 3NaH_2 \stackrel{+1}{PO_2}$$

**118.** (b) 
$$P_4 + NaOH \rightarrow No reaction$$

**120.** (c) 
$$Ca_3P_2 + 6H_2O \rightarrow 3Ca(OH)_2 + 2PH_3$$

122. (b)  $PH_3$  is less basic because lone pair is not easily available for donation.

**123.** (d) 
$$P_2O_3 + 3H_2O \rightarrow 2H_3PO_3$$

**124.** (d) 
$$P_2O_5 + 3H_2O \rightarrow 2H_3PO_4$$
 orthophosphoric acid.

**125.** (c) 
$$H_3PO_2$$
 Monobasic acid

$$\begin{array}{c}
O \\
P \\
P \\
OH
\end{array}$$
 Only one  $-OH$  group

**126.** (b) 
$$PCl_3 + 3H_2O \rightarrow H_3PO_3 + 3HCl$$

**127.** (b) 
$$H_3PO_3$$

$$H \overset{O}{\stackrel{\parallel}{\nearrow} P} OH \\ OH$$

**128.** (c) 
$$H_3 P O_2$$
  
  $3 + x + (-2 \times 2) = 0$   
  $x = +1$ 

**129.** (d)  $Na_4P_2O_7$  Salt of strong acid and strong base.

**130.** (b) 
$$P_4 + 6H_2SO_4 \rightarrow 4H_3PO_4 + 6SO_2$$

**131.** (c) 
$$CaCN_2 + 3H_2O \rightarrow CaCO_3 + 2NH_3$$

**132.** (b) 
$$H_4P_2O_7$$

4 - OH group are present.

**134.** (b) 
$$BiCl_3 + H_2O \rightarrow BiOCl + 2HCl$$

**135.** (b) 
$$CaC_2 + N_2 \xrightarrow{500-600^{\circ}C} CaCN_2 + C$$

**136.** (a) 
$$CaCN_2 + 3H_2O \rightarrow CaCO_3 + 2NH_3$$

**137.** (a) 
$$NH_2CONH_2$$

% of 
$$N = \frac{\text{Mass of } N}{\text{Mass of compound}} \times 100 = \frac{28}{60} \times 100 =$$

46%.

**141.** (a) Anhydride of nitrous acid is  $N_2O_3$ .

**144.** (b) 
$$P_4 + 3NaOH + 3H_2O \rightarrow PH_3 + NaH_2PO_4$$

**145.** (a)  $NH_3$  is highly soluble due to H-bonding.

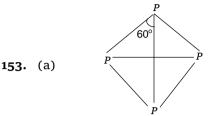
$$\begin{array}{cccc} H & H\text{-bonding} \\ \stackrel{\delta^+}{\sim} & \stackrel{\delta^-}{\sim} & \stackrel{}{\checkmark} & \stackrel{\delta^+}{\sim} & \stackrel{\delta^-}{\sim} \\ H-N...... & H-O \\ & & H & H \end{array}$$

**146.** (d) 
$$NH_3 PH_3 AsH_3 SbH_3$$

B.pt in (K) 238.5 185.5 210.6 254.6

**149.** (a) 
$$H_3PO_2$$
  
  $3+x-4=0$   
  $r=+1$ 

**151.** (d) Solid  $PCl_5$  exists as  $PCl_4^+$  and  $PCl_6^-$ .



**154.** (e) Phosphorus minerals is called as hydroxy apatite and fluorapatite.

**156.** (a) Nitrogen does not have *d*-orbitals.

**157.** (c) 
$$3CuO + 2NH_3 \rightarrow 3Cu + N_2 + 3H_2O$$
.

**159.** (b) Liquid ammonia is used in refrigeration because it has high heat of vaporisation.

**160.** (a) 
$$Sn + conc. 4NHO_3 \rightarrow H_2SnO_3 + 4NO_2 + H_2O_3$$
Meta stannic acid

**161.** (c) 
$$3Cu + 8NHO_3 \rightarrow 3Cu(NO_3)_2 + 4H_2O + 2NO_{\text{Nirric oxid}}$$

163. (c) Pentavalency in phosphorus is more stable that of nitrogen due to large size of phosphorus atom.

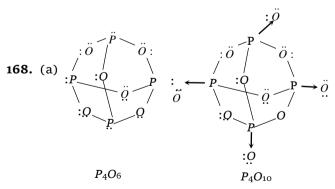
164. (b) Ammomium nitrate is neutral fertilizer.

**165.** (d)  $PH_3$  insoluble in water because does not consist of hydrogen bond.

**166.** (a) 
$$NH_4Cl + NaNO_2 \xrightarrow{\Delta} NH_4NO_2$$
  
 $NH_4NO_2 \xrightarrow{} N_2 + 2H_2O$ 

**167.** (a) Nitrogen react with metal to form a nitride.

$$6Li + N_2 \xrightarrow{450^{o}C} 2Li_3N$$
 (Lithium nitride)



**169.** (b) It is a salt of pyrophosphoric acid  $H_4P_2O_7$ .

**172.** (d) Copper react with conc. nitric acid to form a nitric oxide.

173. (a)  $N_2O$  on account of stimulating effect on nervous system.

174. (d) Sodium metal in liq.  $N\!H_3$  solution shows strong reducing power due to solvated electron.

$$Na + (x + y)NH_3 \Rightarrow [Na(NH_3)_x]^+ + [e(NH_3)_y]^-$$
solvated electron

- **175.** (c)  $PH_3 + 4Cl_2 \rightarrow PCl_5 + 3HCl$
- **178.** (c) Generally  $P_2O_5$  are used as a dehydrating agent.
- **180.** (a) Phosphorus show + 5 valency.
- **181.** (b) In the Haber process for the manufacture of  $NH_3$ , Fe is used catalyst and Mo as a promotre.
- **182.** (a) On adding excess of ammonium hydroxide to a copper chloride solution a deep blue solution of  $[Cu(NH_3)_4]^{2+}$  ion is formed.
- **183.** (d)  $(NH_4)_2 SO_4 + KCNO \rightarrow$

$$NH_4CNO + K_2SO_4 \rightarrow NH_2 - CO - NH_2$$
Urea

- **185.** (a) Nitric acid turns the skin yellow because it reacts with protein giving a yellow compound called xanthoprotein.
- **186.** (d) Ammonium sulphate is a nitrogenous fertilizers.
- **187.** (d) Ammonia generally prepared by the Haber's process.
- **192.** (a)  $H_3PO_2$  is hypophosphorus acid
- **193.** (c)  $(NH_4)_2SO_4$  is a salt of weak base & strong acid

$$(NH_4)_2 SO_4 + 2H_2O \rightarrow 2NH_4OH + H_2SO_4$$
  
Weak base Strong acid

- **194.** (a) One part of concentrated  $HNO_3$  and 3 parts concentrated HCl form aquaregia.
- **196.** (c) -3 to +5  $PH_3(-3)$  and  $H_3PO_4(+5)$
- **199.** (b)  $BiCl_5$  does not exist because +3 oxidation state of Bi s more stable than +5 due to inert pair effect.
- **200.** (c)  $H_3PO_3 \rightarrow \text{Tribasic acid} \rightarrow 3 OH$  groups are present

$$H_3PO_4 \to 3H^+ + PO_4^{3-}$$

$$O$$
 $\parallel$ 
 $P \sim OH$ 

**201.** (d)  $Na_2HPO_4 \rightarrow Na_2PO_4^- + H^+$ 

It can give  $H^+$  ion in solution.

**202.** (c)  $NH_3$  and  $PH_3$  both are basic because of the presence of lone pair of electrons.

203. (b) 
$$\xrightarrow{NH_3 > PH_3 > AsH_3 > SbH_3 > BiH_3}$$

Stability decreases down the group because bond energy decreases down the group.

- **204.** (a) Nitrogen forms  $NH_3$  which is most basic.
- **205.** (b)  $H_3PO_3$  is a diabasic acid. It forms two types of salts  $NaH_2PO_3$  and  $Na_2HPO_3$ .
- **206.** (a)  $NH_2 CO NH_2 + 2HNO_2 \rightarrow CO_2 + 3H_2O + 2N_2$
- **207.** (c) I II III IV V

Element – P As Sb Bi NAtomic no. 15 33 51 83

**210.** (c)  $HO - \stackrel{\parallel}{P} - OH$  it is ionizes in three steps OH

because three -OH group are present.

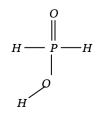
**212.** (c)  $Ca_3P_2 + 3H_2O \rightarrow 3Ca(OH)_2 + 2PH_3$ 

0

- **213.** (d)  $(NH_4)_2Cr_2O_7 \rightarrow N_2 + Cr_2O_3 + 4H_2O_4$
- 214. (b) B > P > As > BiAs we go down the group bond angle

decreases because repulsion between bonded pairs of electron decreases.

- **215.** (b,c)  $3NH_3 + OCl^- \rightarrow NH_2 NH_2 + NH_4Cl + OH^-$
- **217.** (a) Acidic character of oxides decreases down the group.
- **218.** (d)  $N_7 1s^2, 2s^2, 2p^3$  d-orbitals absent in second sub-shell.
- **220.** (c)  $N_2$  can form  $NCl_3$ ,  $N_2O_5$  and  $Ca_3N_2$  but does not form  $NCl_5$ .
- **221.** (a) Highest oxidation state is +5 which remains unchanged.
- **222.** (a) Hypophosphorus acid  $(H_3PO_2)$  is a monobasic acid which act as reducing agent. In this molecule two P-H bonds are responsible for its reducing character and one O-H bond is responsible for its monobasic acid character.
- **223.** (a) Bone black is the polymorphic form of phosphorus.
- 224. (b) Nitrous oxide is known as Laughing gas.
- **225.** (a) We know that,  $2Pb(NO_3)_2 \rightarrow 2PbO + 4NO_2 + O_2$  So nitric oxide  $(NO_2)$  is produced.
- **226.** (d) Phosphorus exist as solid at  $27^{\circ}C$  and 1 atmospheric pressure (m.p. of white phosphorus =  $44^{\circ}C$ )
- **227.** (b) We know that,  $4HNO_3 + P_4O_{10} \rightarrow 4HPO_3 + 2N_2O_5$ The product is dinitrogen pentaoxide  $(N_2O_5)$ .



#### **228.** (b) Hypophosphorous acid is $H_3PO_2$ .

**229.** (b) 
$$NO(g) + NO_2(g) \xrightarrow{-30^{\circ}C} N_2O_3(l)$$
(Blue)

**230.** (c) The ignition temperature of black phosphorus is highest among all allottropes.

**231.** (a) 
$$(NH_4)_2Cr_2O_7 \xrightarrow{\Delta} N_2 \uparrow + Cr_2O_3 + 4H_2O$$
  
 $NH_4NO_2 \xrightarrow{\Delta} N_2 \uparrow + 2H_2O$ 

**232.** (a) Nitrogen shows +I to +V, all oxidation states.

**233.** (c) Boiling points of  $SbH_3(254\ K)$ ,  $NH_3(238\ K)$ ,  $AsH_3(211\ K)$  and  $PH_3(185\ K)$  therefore boiling points are of the order  $SbH_3 > NH_3 > AsH_3 > PH_3$ .

**234.** (a) Because phosphorous is most electronegative element out of P, Bi, Sb and C.

**235.** (d)  $Ca_3(PO_4)_2 + 3SiO_2 \rightarrow 3CaSiO_3 + P_2O_5$ 

**236.** (a) When a solid compound on heating change into gaseous state without changing into liquid state, the phenomenon is known as sublimation. *e.g.*,  $I_2$ ,  $NH_4CI$  and camphor.

**237.** (b) 16 bond by its structure.

**238.** (d) Phosphorus is a non-metallic element. It form's acidic oxide.

**239.** (b) 
$$2\stackrel{+4}{NO}_2 + H_2O \rightarrow HNO_2 + HNO_3$$
 Mixed acid anhydride

**240.** (c) Oxidation number of As in  $H_2AsO_4^-$ 

$$2 + x - 8 = -1$$
$$x - 6 = -1$$
$$x = 5$$

**241.** (a) The inorganic nitrogen exists in the form of ammonia, which may be lost as gas to the atmosphere, may be acted upon by nitrifying bacteria, or may be taken up directly by plants.

## Oxygen family

- (c) Sulphur -(1) Monoclinic (2) Rhombic (3) Plastic
- **4.** (c)  $S + O_2 \rightarrow SO_2$

5. (a) 
$$\frac{1}{8}S_8 + 2e^- \rightarrow S^{2-}$$

**6.** (c)  $2H_2O + 2F_2 \rightarrow 4HF + O_2$ 

9. (b) 
$$O_8$$
 -  $1s^2$   $2s^2$   $2p^4$ 

1 1 2 1 1 2 2 unpaired electrons

**11.** (a) Element – O S Sc Te Po

13. (b,c)  $Na_2SO_3 + H_2SO_4 \rightarrow Na_2SO_4 + H_2O + SO_2$ (S) (aq) (d) (g)

**15.** (d)  $H_2O$   $H_2S$   $H_2Se$   $H_2Te$   $104.5^{\circ}$   $92.1^{\circ}$   $91^{\circ}$   $90^{\circ}$ 

As we go down the group electronegativity decreases due to which repulsion between bonded pairs of electron also decreases. Hence, bond angle decreases.

**20.** (a) Paramagnetism because of two unpaired electrons in the antibonding molecular orbitals.

**21.** (a)  $2Na_2SO_3 + O_2 \rightarrow 2Na_2SO_4$ 

**24.** (c) 
$$3O_{\frac{\text{silent}}{\text{electric discharge}}} 2O_3$$

**28.** (b) 
$$O_3 \rightarrow O_2 + [O]$$

$$2KI + H_2O + [O] \rightarrow 2KOH + I_2$$

$$2KI + H_2O + O_3 \rightarrow 2KOH + I_2 + O_2$$

**32.** (d)  $2KMnO_4 + 3H_2SO_4 + 5H_2S \rightarrow$ 

$$K_2SO_4 + 2MnSO_4 + 8H_2O + 5S$$

**33.** (a)  $Cu + 2H_2SO_4 \rightarrow CuSO_4 + 2H_2O + SO_2$ 

**35.** (a) The minimum and maximum oxidation number of S are -2 and +6 respectively. Since the oxidation number of S in  $SO_2$  is +4, therefore it can be either increased or decreased. Therefore  $SO_2$  behaves both as an oxidising as well as reducing agent.

**36.** (a)  $2H_2S + SO_2 \rightarrow 2H_2O + 3S$  reducing agent oxidising agent

37. (d)  $K_2Cr_2O_7 + H_2SO_4 + 3SO_2 \rightarrow$ 

$$K_2SO_4 + Cr_2(SO_4)_3 + 3H_2O$$
Green

**39.** (a)  $SO_2 + 2MgO + S$ 

**40.** (a)  $2H_2O + SO_2 \rightarrow H_2SO_4 + 2[H]$  (nascent hydrogen) Coloured flower  $+2[H] \rightarrow$  Colourless flower

**41.** (c) 
$$H_2SO_3 + 2NaOH \rightarrow Na_2SO_3 + 2H_2O$$
Sodium sulphite

**42.** (a)  $H_2O + SO_3 \rightarrow H_2SO_4$ 

**43.** (d) 
$$H_2SO_4 + SO_3 \rightarrow H_2S_2O_7$$

**45.** (b)  $2Ag + 2H_2SO_4 \rightarrow Ag_2SO_4 + 2H_2O + SO_2$ Reducing Agent agent agent

**46.** (a) Only dehydrating agent  $HCOOH \xrightarrow{H_2SO_4} CO + H_2O$ 

- **48.** (b)  $COOH \xrightarrow{\text{conc.} H_2SO_4} H_2O + CO + CO_2$
- **51.** (d)  $H_2SO_4 + SO_3 \rightarrow H_2S_2O_7$  (Oleum or Furning  $H_2SO_4$ )
- **53.** (d)  $S_2 O_7^{2-}$

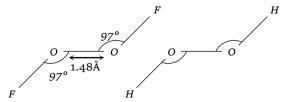
- **55.** (d)  $Na_2SO_3 + S \rightarrow Na_2S_2O_3$
- **56.** (b)  $Na_2S_2O_3 + Cl_2 + H_2O \rightarrow Na_2SO_4 + 2HCl + S$
- **58.** (c)  $AgBr + 2Na_2S_2O_3 \rightarrow Na_3[Ag(S_2O_3)_2] + NaBr$
- **60.** (a)  $SO_2 + 2H_2S \rightarrow 3S + 2H_2O$
- **63.** (a)  $2H_2S + O_2 \rightarrow 2H_2O + 2S$
- **64.** (a) Mixture of  $K_2Cr_2O_7$  and conc.  $H_2SO_4$  is known as chromic acid.
- **65.** (c)  $H_2O$  contain hydrogen bond while no hydrogen bonding is present in  $H_2S$ .
- **67.** (a)  $H_2O$  containing hydrogen bond.
- **69.** (c)  $HO SO_2 OH + PCl_5 \rightarrow Cl SO_2 OH + POCl_3$

+HCl

$$HO-SO_2-OH+2PCl_5 \rightarrow$$

$$Cl - SO_2 - Cl + 2POCl_3 + 2HCl$$
  
Sulphuryl chloride

- **70.** (d) Mixture of  $O_2$  and  $CO_2$ .
- **71.** (c) The bond between  $(H_2Te)$  is weakest hence it gives  $H^+$  ion easily.
- **73.** (c)  $KO_2$  because in  $O_2^-$  (superoxide ion) One unpaired electron is present in the antibonding orbital.
- **75.** (b) Oxidation states are 2, 4, 6
- 77. (b)  $H_2O$   $H_2S$   $H_2Se$   $H_2Te$  373K 213K 269K 232K  $H_2S$  has lowest boiling point and  $H_2O$  has highest boiling point because if any compound has hydrogen bond. Its boiling point is high.
- **78.** (c)  $V_2O_5$  catalyst are used for the manufacture of  $H_2SO_4$  by Contact process.
- **81.** (d) *O*, *S*, *Se* shows polymorphism.
- **84.** (e)  $H_2S + 2HNO_3 \rightarrow 2NO_2 + S + 2H_2O$ .
- **85.** (b)  $O_2F_2$  is similar to that of  $H_2O_2$



- **87.** (d) 16<sup>th</sup> group called chalcogens (oxygen family) while *Na* is a I<sup>st</sup> group element which is called alkali metal.
- **90.** (b) Caro's acid  $(H_2SO_5)$
- **91.** (c)  $2KO_2 + CO_2 \rightarrow K_2CO_3 + \frac{3}{2}O_2$
- **93.** (a)  $H_2O$  consist of highest boiling point than other hydride (Due to presence of the hydrogen bonding).
- 94. (b) Lose of electron.
- **95.** (c)  $SO_2$  is soluble in water

$$\begin{array}{c} H_2O + SO_{\,2} \,\, \rightarrow \,\, H_2SO_{\,3} \\ \qquad \qquad \text{Sulphurus acid} \end{array}$$

- **96.** (c)  $2Na_2O_2 + 2H_2O \rightarrow 4NaOH + O_2$ . In this reaction oxygen  $(O_2)$  is formed.
- **97.** (b)  $3S + 4NaOH \xrightarrow{\text{boiling}} Na_2S_2O_3 + Na_2S$
- **98.** (c) Quartz is a crystalline variety of silica.
- **99.** (b) 98%  $H_2SO_4$  is used for absorbing dense fog of acid which is formed by dissolving  $SO_3$  in water. Hence 98%  $H_2SO_4$  is the most efficient agent for the absorption of  $SO_3$ .
- **100.** (b) Concentrated  $H_2SO_4$  is diluted by adding the conc.  $H_2SO_4$  in the water drop by drop with constant stirring because it is an exothermic reaction and by doing so heat is generated slowly and dissipated in the atmosphere.
- **101.** (a) Photochemical smog is caused by oxides of sulphur and nitrogen.
- **102.** (d)  $SO_2$  act as bleaching agent due to its reducing property.

$$SO_2 + 2H_2O \rightarrow H_2SO_4 + 2H$$

Coloured matter  $+H \rightarrow$  Colourless matter

- 103. (b) Ozone on treatment with  $SO_2$  produce  $SO_3$ .  $3SO_2 + O_3 \rightarrow 3SO_3$
- **104.** (b)  $PbO_2$  is a powerful oxidizing agent and liberate  $O_2$  when treated with acids SO,  $O_2$  gas will be evolved.
- **105.** (c) Dithionous acid  $(H_2S_2O_4)$  has sulphur in +3 oxidation state.

$$O O O HO - S - S - OH$$
;  $2(+1) + 2x + 4(-2) = 0$   
 $2x = 8 - 2 = 6$ ;  $x = +3$ 

- **106.** (a)  $SO_2$  acts as an oxidising agent particularly when treated with stronger reducing agents.  $SO_2$  oxidises  $H_2S$  into S.
- **107.** (a)  $R_3SiCl$  on hydrolysis forms only a dimer.  $R_3SiOH + HOSiR_3 \rightarrow R_3Si O SiR_3$ .

108. (d) Formation of chlorine nitrate is the main cause of ozone depliction.

# Halogen family

- (b) HF > HCl > HBr > HI (Thermal stability). 1.
- (a)  $CHCl_3 + \frac{1}{2}O_2 \rightarrow {COCl_2 + HCl}_{\begin{subarray}{c} Phosgene or carbonyl chloride \end{subarray}}$ 2.
- (d) Iodine has the least affinity for water and is 3. only slightly soluble in it. However, it dissolves in 10% aq. solution of KI due to the formation of a complex ion *i.e.*  $I_3^-$ .

$$I_2 + KI = KI_3$$
 or  $I_2 + \Gamma = I_3^-$  (complex ion)

- (c)  $2Na_2S_2O_3 + I_2 \rightarrow 2NaI + Na_2S_4O_6$ 4.
- (a)  $Cl_2 + 2KBr \rightarrow 2KCl + Br_2$ 5.

A more electronegative halogen can displaces less electronegative halogen.

- (a) HI is the strongest reducing agent among 6. halogen acids because of lowest bond dissociation energy.
- 8. (a) Due to H-Bonding free ions are not present in aq. solution. Hence, bad conductor.
- (c) Electronegativity of  $I_2$  is less than  $Br_2$ . 9. Therefore unable to displace bromine.
- (b) Carnellite is  $KCl.MgCl_2.6H_2O$ . The mother 10. liquor life after crystallisation of KCl from carnellite contain about 0.25% of bromine as  $MqBr_2$  and KBr.
- (a) HF is liquid because of intermolecular H-11. Bonding.
- (a)  $\frac{HClO}{\text{Weak acid}} = H^+ + \frac{ClO}{\text{Strong conjugate}}$ 12.
- (d)  $2NaOH + Cl_2 \xrightarrow{\text{Cold}} NaCl + NaClO + H_2O$ 13.  $6NaOH + 3Cl_2 \xrightarrow{\text{heat}} 5NaCl + NaClO_3 + 3H_2O$
- (b)  $6KOH + 3Cl_2 \rightarrow 5KCl + KClO_3 + 3H_2O$ . 15.
- 17. (a) HF is the weakest acid. Since it is unable to give  $H^+$  ions which are trapped in H-Bonding.
- (b) Hydride -HFHCl HBrHI 18. B.pt (in K) - 293 189 206 238 Becouse of having low b.p. HCl is more volatile.
- (a)  $2KClO_3 + I_2 \rightarrow 2KIO_3 + Cl_2$ 19.
- (c)  $2Na_2S_2O_3 + I_2 \rightarrow 2NaI + Na_2S_4O_6$ 20.
- (d)  $2KMnO_4 + 16HCl \rightarrow 2KCl + 2MnCl_2 + 5Cl_2 + 8H_2O$ 21.
- +5 (a) 22. +3  $HClO_4 > HClO_3 > HClO_2 > HClO$

As the oxidation no. of halogen increases acidic character increases.

(c)  $2KBr + 3H_2SO_4 + MnO_2 \xrightarrow{\Delta}$ 

$$2KHSO_4 + MnSO_4 + 2H_2O + Br_2$$

(b,d) Electron affinity of  $Cl_2$  is maximum 29.

Boiling 185.2 pt -34.659.5  $(^{\circ}C)$ 188.1

- **32.** (a)  $Cl_2 + 2NaBr \rightarrow 2NaCl + Br_2$
- 33. (d)  $CCl_4 + H_2O \rightarrow No reaction$ d-orbitals are absent in carbon atom.
- (a)  $I_2 + 10HNO_3 \rightarrow 2HIO_3 + 10NO_2 + 4H_2O$ 34.
- (d)  $KI + I_2 \rightarrow KI_3$ 35.
- (a)  $2KBr + H_2SO_4 \rightarrow K_2SO_4 + 2HBr$ 36.
- (b)  $H_2 + F_2 \to 2HF$ 37.
- (b)  $CuSO_4 + 2KI \rightarrow CuI_2 + K_2SO_4$  $2CuI_2 \rightarrow 2CuI$ Cuprous iodide
- atomic number (d) As the increases 43. electronegativity decreases. Hence, tendency to gain electron decreases.
- **51.** (a)  $F_2 + 2Cl^- \rightarrow Cl_2 + 2F^ F_2 + 2Br^- \rightarrow Br_2 + 2F^ F_2 + 2I^- \rightarrow I_2 + 2F^-$
- **53.** (d)  $Br_2 + 2KI \rightarrow I_2 + 2KBr$
- **56.** (d)  $2F_2 + 2H_2O \rightarrow 4HF + O_2$  $3F_2 + 3H_2O \rightarrow 6HF + O_3$
- (c)  $2NaCl + 2H_2O \rightarrow 2NaOH + Cl_2 + H_2$ (anode) (cathode)
- 59.
- (b)  $2NaCl + 2H_2O \xrightarrow{\text{Electroly is}} 2NaOH + Cl_2 + H_2$
- (c)  $H_2O + Cl_2 \rightarrow HCl + HClO$ 62.  $HClO \rightarrow HCl + [O]$  or  $2HClO \rightarrow 2HCl + O_2$
- **63.** (a)  $2NaOH + Cl_2 \rightarrow NaClO + NaCl + H_2O$
- (b)  $Cl_2 + H_2O \rightarrow 2HCl + [O]$  Nascent oxygen Coloured  $+[O] \xrightarrow{\text{Bleaching}} \text{Colourless flower}$ agent flower
- **68.** (b)  $CaO + Cl_2 \rightarrow CaOCl_2$  $NaHCO_3 + Cl_2 \rightarrow No reaction$
- (c)  $Ca(OH)_2 + Cl_2 \rightarrow CaOCl_2 + H_2O$ 69.
- 71. (b)  $MgBr_2 + Cl_2 \rightarrow MgCl_2 + Br_2$

**72.** (a) 
$$Cl_2 + 2Br^- \rightarrow 2Cl^- + Br_2$$

73. (a) 
$$KI + H_2SO_4 \xrightarrow{\Delta} KHSO_4 + HI$$
Conc.

Conc. 
$$H_2SO_4 + 2HI \rightarrow 2H_2O + I_2 + SO_2 \uparrow$$
  
Violet vapour

**74.** (b) 
$$2NaI + Cl_2 \rightarrow 2KCl + I_2$$
 
$$CCl_4 + I_2 \rightarrow \text{Violet colour}$$

76. (b) 
$$KI + H_2SO_4 \xrightarrow{\Delta} KHSO_4 + HI$$
  
 $H_2SO_4 + 2HI \rightarrow 2H_2O + I_2 + SO_2 \uparrow$   
Violet vapour

**79.** (c) 
$$KI + H_2SO_4 \rightarrow KHSO_4 + HI$$

HI is formed but it is further oxidised by conc.  $H_2SO_4$  into  $I_2$ 

$$2HI + H_2SO_4 \rightarrow 2H_2O + I_2 + SO_2 \uparrow$$
Violet vapour

**80.** (b) 
$$HCl \xrightarrow{H_2O} H_{(aq)}^+ + Cl_{(aq)}^-$$

**81.** (b) 
$$4NaCl + K_2Cr_2O_7 + 3H_2SO_4 \rightarrow K_2SO_4 + 2Na_2SO_4 + 2CrO_2Cl_2 + 3H_2O_4$$
Chromylchloride

**82.** (c) Hydrogen bonding is absent in 
$$HI$$
 while it is present in  $NH_3, H_2O$  and  $C_2H_5OH$ .

**85.** (d) 
$$HI$$
 is strongest acid because  $H-I$  bond is weakest among halogen acids.

**88.** (a) 
$$ClO_2^-$$
 has  $sp^3$ -hybridization and two lone pairs on halogen which produces V-shape Bent structure

**89.** (d) 
$$2HClO_4 \rightarrow H_2O + Cl_2O_7$$
.

**90.** (b) 
$$F_2$$
 gases  $Cl_2$   $Br_2$  liquid  $I_2$  solid

As we go down the group Vander Waal forces increases. Hence, physical state changes.

**92.** (c) *F* can not act as reducing agent because it has highest reduction potential

$$F_2 + 2e^- \rightarrow 2F^-; E^o = +2.87 V$$

**93.** (c) 
$$I_2 + 10HNO_3 \rightarrow 2HIO_3 + 10NO_2 + 4H_2O_3$$

**94.** (a) 
$$IF_5 + F_2 \rightarrow IF_7$$
.

Pseudohalide Pseudohalogens  $CN^-$  cyanide  $(CN)_2$  Cyanogen

SCN - Thiocyanate (SCN)<sub>2</sub> Thiocyanogen

**96.** (c) *NaF* is highest melting halide because it is most ionic in nature.

**98.** (b) 
$$2F_2 + 2H_2O \rightarrow 4HF + O_2$$

halogens.

**99.** (c) 
$$CaOCl_2 \rightarrow CaCl_2 + [O]$$
 Nascent oxygen

**100.** (a) Generally alkali metals and alkali earth metals elements extracted by the fused electrolysis method.

**102.** (c) 
$$2Na_2S_2O_3 + I_2 \rightarrow 2NaI + Na_2S_4O_6$$
.

104. (b) Beilstein test – In this test organic compound is heated on a copper wire in a flame. The appearance of a green or bluish green flame due to the formation of volatile cupric halides indicate the presence of halogens in the organic compound. (It does not tell which halogen is actually present).

**105.** (d) 
$$Cl_2 + 2KBr \rightarrow Br_2 + 2KCl$$
 Oxidation

**107.** (b) 
$$3HCl + HNO_3 \rightarrow NOCl + 2H_2O + Cl_2$$
.

**108.** (a) 
$$Cl - Cl \xrightarrow{\text{U.V}} Cl + Cl$$
Free radical

**109.** (a) *HF* is a weak acid due to intermolecular hydrogen bonding.

**110.** (a) Acidic nature of oxide ∞ Non metallic nature of element

Non metallic nature decrease in the order Cl > S > P.

111. (c) Aqua regia is 1 part of  $HNO_3$  and 3 part of HCl.

**113.** (a) *AqI* is a covalent compound.

114. (a) Bromine is a liquid at room temperature.

**115.** (a) 
$$Cl_2 + H_2O \rightarrow 2HCl + [O]$$
Nascent oxyge

**117.** (b) The enamel of our teeth is the hardest substance in the body made up of  $CaF_2$  and dentine below it made of  $Ca_3(PO_4)_2$ .

- **118.** (b) As the electronegativity decreases reactivity also decreases.
- **119.** (b)  $KI + I_2 \rightarrow KI_3$  (soluble complex)
- **121.** (d)  $HI < I_2 < ICl < HIO_4$
- **122.** (a) HF < HCl < HBr < HCl

As we go down the group bond energy decreases hence, acidic nature increases.

- **123.** (b) Caliche is crude chile salt petre  $(NaNO_3)$  which contains about 0.02% iodine as sodium iodate  $(NaIO_3)$ .
- **124.** (a) *LiF > LiCl > LiBr > LiI*Lattice energy depends on the size and charge of the ion.
- **125.** (a) *F-F* more strong bond compare to *F-Cl*, *F-Br* and *Cl-Br* bond.
- **126.** (c)  $2Na_2S_2O_3 + I_2 \rightarrow Na_2S_4O_6 + 2NaI$ .
- **127.** (d)  $CaCl(OCl) \Rightarrow Ca \stackrel{\checkmark}{\searrow} Cl$
- **128.** (a)  $2NaOH + Cl_2 \rightarrow NaCl + NaOCl + H_2O$  $\downarrow \qquad \qquad \downarrow \qquad \downarrow \qquad \qquad$
- 131. (d)  $I_2 + NaF \longrightarrow$   $I_2 + NaBr \longrightarrow$   $I_2 + NaCl \longrightarrow$ No reaction

Because  $I_2$  is least electronegative among halogens.

- **132.** (a)  $HClO_4 > HCl > H_2SO_4 > HNO_3$ Decreasing order of acidic character.
- **134.** (a)  $HgCl_2 + Hg(CN)_2 \rightarrow HgCl_2 . Hg(CN)_2$ Mercuric chloride Mercuric cyanide
- **135.** (a) HI > HBr > HCl > HF Acidic character decreasing order.
- 136. (c)  $Ca(OH)_2 + Cl_2 \rightarrow CaOCl_2 + HOCl$ Slaked lime Bleaching poeder
- **137.** (d)  $K_2Cr_2O_7 + 14HCl \rightarrow 2KCl + 2CrCl_3 + 7H_2O + 3Cl_2$
- **139.** (d) Fluorine does not gives positive oxidation state it is always show -1 oxidation state.
- **140.** (a)  $HClO_4 > HClO_3 > HClO_2 > HCl$
- **141.** (d)  $Ca(OH)_2 + Cl_2 \rightarrow CaOCl_2 + H_2O$  bleaching powder
- **146.** (a)  $F_2 + (dil.) 2NaOH \rightarrow 2NaF + OF_2 + H_2O$
- **147.** (a) Fluorine can not be oxidised by even strongest oxidising agent.
- **148.** (b)  $Br_2 + 2H_2O + SO_2 \rightarrow H_2SO_4 + 2HBr$
- 149. (c)  $2KBr + I_2 \rightarrow 2KI + Br_2$ Iodine is a less electronegative compare to Bromine hence iodine does not change  $Br^-$  to  $Br_2$ .
- **150.** (a)  $CaI_2$  are show covalent properties than other  $CaF_2$ ,  $CaCl_2$ ,  $CaBr_2$  compound.

**151.** (d)  $2KMnO_4 + 3H_2SO_4 + 10HCl \rightarrow$ 

$$K_2SO_4 + 2MnSO_4 + 8H_2O + 5Cl_2$$

- 153. (d)  $PbI_4$  is least stable because of two reasons
  - (1) Size of iodine is biggest.
  - (2) +2 oxidation state of Pb is more stable than +4 state because of inert pair effect.
- **154.** (a)  $Cl_2 + NaF \rightarrow No reaction$

Since  $\operatorname{Cl}_2$  is less electronegative then  $\operatorname{F}_2$ . Therefore unable to displace fluorine from its salt.

- **156.** (b)  $CS_2 + 3Cl_2 \xrightarrow{l_2} CCl_4 + S_2Cl_2$
- **157.** (b) According to the Fajan's rule largest cation and smallest anion.
- **158.** (a)  $2F_2 + 4NaOH \rightarrow 4HF + 2H_2O + O_2$
- **159.** (c)  $Cl_2 \rightarrow 2Cl \Delta H = +ve$ 1 mole 2 moles

  High temperature and low pressure is favourable.
- **160.** (d)  $BF_3$  accept lone pair of electrons.
- **161.** (d)  $CrO_2Cl_2$  is a orange red gas.
- 164. (a) Florine always show -1 oxidation state.
- **165.** (a) Solid *NaF* is used to purify fluorine *i.e.* by removing of *HF* fumes.
- **166.** (c)  $KHF_2 \rightarrow KF + HF$

$$KF \rightarrow K^+ + F^-$$

At cathode :  $K^+ + e^- \rightarrow K$ 

$$2K + 2HF \rightarrow 2KF + H_2$$

At anode:  $F^- \rightarrow F + e^-$ 

$$F+F \rightarrow F_2$$

- **168.** (c) Small atomic size of *Li* and *F* lattice energy is highest.
- **169.** (b)  $SO_2$  bleaches flower by reduction

$$2H_2O + SO_2 \rightarrow H_2SO_4 + 2[H]$$

$$2[H] + Coloured \xrightarrow{\text{Reduction}} Colourless$$
 flower reduced flower

This bleaching is temporary because reduced flower again oxidised by air to form coloured flower

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

$$\begin{array}{c} [O] + \text{Coloured} & \xrightarrow{\text{Oxidation}} & \text{Colourless} \\ \text{flower} & \text{Oxidised flower} \end{array}$$

This bleaching is permanent because oxidised flower remains colourless.

- **170.** (a) Fluorine does not form oxyacids because it is more electronegative than oxygen.
- **173.** (c)  $MnO_2 + 4HCl \rightarrow MnCl_2 + 2H_2O + Cl_2$

$$2KMnO_4 + 3H_2SO_4 + 10HCl \rightarrow$$

 $K_2SO_4 + 2MnSO_4 + 8H_2O + 5Cl_2$ 

175. (b) 
$$3KClO_3 + 3H_2SO_4 \xrightarrow{\text{Heat}}$$

$$3KHSO_4 + HClO_4 + 2ClO_2 + H_2O$$

- **176.** (b) F > Cl > Br > I. As the size increases electronegativity decreases.
- **177.** (c) Ionic radius increases on going down the group because no. of shells increases.
- **178.** (b) Reducing properties increase from F to I so it oxidise by nitric acid.  $I_2 + 10 HNO_3 \rightarrow 2 HIO_3 + 10 NO_2 + 4 H_2 O$
- **179.** (a) Fluorine and chlorine are more electronegative than sulphur.
- **180.** (d) Upper halogen can replace lower halogen from their compounds solution.
- **181.** (a) Iodine  $(I_2)$  is slightly soluble in water but it dissolves in 10% aqueous solution of *KI* due to the formation of potassium triodide  $(KI_3)$ .
- **182.** (a) Due to highest electronegativity of fluorine the anion [F--H-F] exists as a result of strong hydrogen bond by which  $K^+$  associate to form  $KHF_2$ .
- **183.** (a) Fluorine is the most electronegative element. It does not form oxyfluoxides like other halogens. If reacts with *NaOH* to form sodium fluoride and oxygen fluoride.  $2NaOH + 2F_2 \rightarrow 2NaF + OF_2 + H_2O$
- **184.** (c) Due to unpaired  $e^ ClO_2$  is paramagnetic.
- **185.** (a) Oxidation number of  $HBrO_4$  is more than that of HOCl,  $HNO_2$  and  $H_3PO_3$  so it is the strongest acid among these.
- **186.** (a) Chlorine heptachloride  $(Cl_2O_7)$  is the anhydride of perchloric acid.  $2HClO_4 \stackrel{\Delta}{\longrightarrow} Cl_2O_7 + H_2O$
- **187.** (c)  $I_2$  forms complex ion  $I_3^-$  in KI solution due to which it dissolves in it.

#### Noble gases

- 3. (a) Gas  $H_2$   $O_2$  He  $N_2$ Mol. mass - 2 32 4
- 5. (c) Helium (In Greek Helios = Sun)
- **6.** (d) All the noble gases are monoatomic, colourless and odourless gases. Their monoatomic nature is due to the stable outer configuration  $ns^2np^6$  of their atoms. As a result, they do not enter into chemical combination even amongst themselves.
- 7. (c) Except *He*, all other noble gases are adsorbed by coconut charcoal at low temperatures. The extent of adsorption increases as the atomic size of the noble gas increases.

- 10. (b) An oxygen-helium mixture is used artificial respiration in deep sea diving instead of air because nitrogen present in air dissolves in blood under high pressure when sea diver goes into deep sea. When he comes to the surface, nitrogen bubbles out of the blood due to decrease in pressure, causing pains. This disease is called "bends".
- 11. (c)  $XeF_2$ ,  $XeOF_2$ .  $XeF_4$ ,  $XeOF_4$ .  $XeF_6$ ,  $XeO_3$ .
- **12.** (c) **Gas** (Abundance in air by volume (ppm))

Helium	5.2
Neon	18.2
Argon	93.4
Krypton	1.1
Xenon	0.09

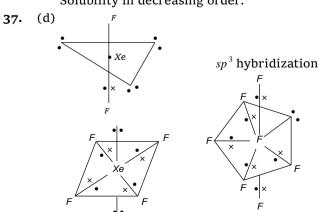
- 13. (c) Neon  $\rightarrow$  Ne is monoatomic and others are diatomic  $N_2, F_2$  and  $O_2$ .
- **14.** (c)  $_1H^2 +_1 H^2 \rightarrow _2He^4$
- **15.** (b)  $HeF_4$  does not exist.
- **16.** (d)  $Ar_{18} \rightarrow 2, 8, 8$
- 17. (b)  $Ne_{10} \rightarrow 1s^2 2s^2 2p^6$
- **23.** (d)  $XeF_2$  has  $sp^3d$  -hybridization with linear shape

$$F$$
 $Xe$ 
 $F$ 

- **24.** (b) Partial hydrolysis;  $XeF_4 + H_2O \rightarrow XeOF_2 + 2HF$ Complete hydrolysis;  $2XeF_4 + 3H_2O \rightarrow Xe + XeO_3 + F_2 + 6HF$
- **26.** (d) *He* is least polarizable because of small atomic size
- 27. (a) Rn because it is radioactive element obtained by the disintegration of radium  ${}_{88}Ra^{206} \rightarrow {}_{86}Rn^{202} + {}_{2}He^{4}$

**30.** (c) 
$$1s^2 2s^2 2p^6 \rightarrow \text{Neon}$$

- **35.** (a)  $XeF_6 + 3H_2O \rightarrow XeO_3 + 6HF$
- **36.** (c) Xe > Kr > Ar > Ne > He Solubility in decreasing order.

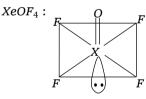


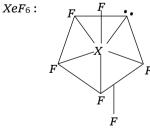
 $sp^3d^2$  hybridization  $sp^3d^3$  hybridization

- (b) Zero group element are show less chemically activity because this group element have 8 electron.
- 39. (d) Xe is formed following compounds.  $XeF_2$ ,  $XeF_4$ ,  $XeF_6$ .
- (a) As the number of shells increases, size 40. increases and the effective nuclear charge on the outermost electron decreases. Thus, I.E. decreases.
- (c)  $XeF_2$ ,  $XeF_4$  &  $XeF_6$  can be directly prepared 44.  $Xe + F_2 \xrightarrow{Ni \text{ tube}} XeF_2$ ;  $Xe + 2F_2 \xrightarrow{673 K} XeF_4$  $Xe + 3F_2 \xrightarrow{523-573 K} XeF_6$

 $XeO_3$  is obtained by the hydrolysis of  $XeF_6$  $XeF_6 + 3H_2O \rightarrow XeO_3 + 6HF$ 

- (a)  $XeO_3$  shows  $sp^3$  hybridization. 46.
- (a) It is because 47. (1) Small atomic size (2) High Ionization energy (3) Absence of *d*-orbitals
- (a) Zero group element are attached with weak 48. intermolecular force.
- 49. (b)  $XeF_2$ ,  $XeF_4$ ,  $XeF_6$ .
- (d) XeO<sub>3</sub>: 50.





- (d) Neil Bartlett prepared first noble gas 51. compound. Xenon hexafluoroplatinate (IV).
- (d) He, Ne, and Kr all are found in very little 53. amount in atmosphere, so all are called rare
- (c) Helium is twice as heavy as hydrogen, its 54. lifting power is 92% of that of hydrogen. Helium has the lowest melting and boiling

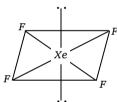
point of any element which makes liquid helium an ideal coolant for many extremely low temperature application such as super conducting magnet and cryogenic research where temperature close to absolute zero are needed.

(a) The maximum temperature at which gas can 55. be liquified is called its critical temperature. The gas which have high boiling point will change into liquid first and so critical temperature of the gas will be more.  $T_C \propto \text{B.P.}$  and B.P.  $\propto \text{Molecular weight}$ So *Kr* liquifies first.

**56.** (c) Suppose the oxidation state of Xe is x.  $XeOF_{2}$ 

 $x + (-2) + 2(-1) = 0 \implies x - 2 - 2 = 0 \implies x = 4$ .

- He Ne **57.** (a) Ar Kr Хe Boiling point of - 269 -246 - 186 -153.6 108.1 -62 Inert gases
- (d) Xe is highly polar since the ionisation 59. potential of xenon is quite close to the ionisation potential of oxygen.
- **60.** (d) In the formation of  $XeF_4$ ,  $sp^3d^2$  hybridisation occurs which gives the molecule an octahedral structure. The xenon and four fluorine atoms are coplanar while the two equatorial positions are occupied by the two lone pairs of electrons.



# **Critical Thinking Questions**

- (c) Correct sequence is  $NH_2CONH_2 > NH_4N_3 > NH_3NO_3 > NH_4Cl$ .
- (d) Second group elements are show strong 2. reducing properties but less Ist group element show less.
- (a) The size of alkaline earth metals is smaller as 3. compared to its corresponding alkali metals and its effective nuclear charge is also more than that of its corresponding alkali metals.
- (d) Lead is maximum in flint glass. 4.
- (b)  $BaSO_4 + 4C \xrightarrow{\text{Heat}} BaS + 4CO$ 5. Thus, on heating they produce BaS + 4CO.
- (d) Smaller the atomic size tendency of hydration 6. is more as the size increases tendency for hydration also decreases.
- (a) Fusion mixture is  $Na_2CO_3 + K_2CO_3$ . 7.
- 8. (b) HCl is a gas.
- (a) (A) Peroxide is  $H_2O_2(4)$ ; (B) Super oxide is  $KO_2(3)$

- (C) Dioxide is  $PbO_2(2)$ ; (D) Suboxide is  $C_3O_2(1)$
- 10. (a)  $H_2SO_4 + SO_3 \rightarrow H_2S_2O_7$ 98% Oleum (Fuming  $H_2SO_4$
- 11. (c)  $SnO_2 + 2NaOH \rightarrow Na_2SnO_3 + H_2O$  $SnO_2 + 4HCl \rightarrow SnCl_4 + 2H_2O$
- **12.** (b)  $NaOH + Al^{+3} \rightarrow NaAlO_2$  Sod. metaluminate (Soluble in NaOH)

 $NaOH + Fe^{+3} \rightarrow No$  reaction (Insoluble in NaOH)

- 13. (d) The composition of the common glass is  $Na_2O.CaO.6SiO_2$  .
- 14. (c) Lead form nitric oxide with dil.  $HNO_3$  $3Pb + 8HNO_3 \rightarrow 3Pb(NO_3)_2 + 2NO + 4H_2O$
- 15. (b) VF

  Xe

  Therefore, total number of lone pair of electron on central atom *Xenon* =1
- **16.** (d) Order of acidic strength is  $H_2Te > H_2Se > H_2S > H_2O$

 $Na_2O$  is a salt of  $NaOH + H_2O$  and  $H_2O$  is least acidic among given acids hence pH in this case will be maximum.

- 17. (a)  $HNO_3$  is the strong oxidising acid so it react with alkali while rest can be react with both and alkali.
- **18.** (d) Zero group is called a buffer group because it lies between highly electronegative halogens and highly electropositive alkali metal element.
- **19.** (c)  $PbSO_4$  is insoluble in water.
- **20.** (a)  $N_2O_5$  is most acidic.
- **21.** (b) Element Fluorine Chlorine Bromine Iodine B.E. in  $kJ \mod e^{-1}$  158.8 242.6 192.8 151.1
- 22. (d)  $CaCN_2 + 3H_2O \xrightarrow{\text{under}} CaCO_3 + 2NH_3$ .
- 23. (a)  $6O_2 \xrightarrow{\text{Ozonoly sis}} 4O_3$

Complete ozonalysis gives 4 volume of ozone.

- **24.** (b) Marble statue  $\rightarrow CaCO_3$ ; Calcinated Gypsum  $\rightarrow CaSO_4.2H_2O$ ; Sea shell  $\rightarrow CaCO_3$ ; Dalomite  $\rightarrow CaCO_3.MgCO_3$ .
- **25.** (b) Sodium is basic in nature.
- **26.** (c)  $I_2 + I^- \rightarrow I_3^-$
- **27.** (a) The hydration energy decreases from  $Be^{+2}$  to  $Ba^{+2}$ . This causes the decrease in the

solubilities of the sulphates in this order. In other words, the solubilities decreases with increase in the ionic size.

- **28.** (d)  $2Mg + O_2 \rightarrow 2MgO$  $3Mg + N_2 \rightarrow Mg_3N_2$
- **29.** (a)  $ZnO + BaO \xrightarrow{1100^{\circ}C} BaZnO_{2}$
- **30.** (b) There are no free electron in  $N_2O_4$ , so it is dimagnetic in nature.
- 31. (b)  $CaSO_4 \cdot 2H_2O \xrightarrow{120^{\circ}C} CaSO_4 \cdot \frac{1}{2}H_2O + 1\frac{1}{2}H_2O$ Ploster of paris
- **32.** (d) Third alkaline earth metal is calcium  $_{20}$   $Ca^{40}$ . No. of electron are 20 & No. of proton are 20; e/20, p/20.
- **33.** (a) In the compounds of alkaline earth metals all the electrons are paired. Hence, they are diamagnetic in nature.
- **34.** (a) Mixture of helium and oxygen is used in artificial respiration.
- **35.** (a) Alkaline solution of pyrogallol quickly absorbs oxygen.
- **36.** (d)  $NH_4NO_2 \rightarrow N_2 + 2H_2O$   $2NaN_3 \rightarrow 2Na + 3N_2$  $(NH_4)_2Cr_2O_7 \rightarrow N_2 + Cr_2O_3 + 4H_2O$
- **37.** (b) Because yellow phosphorus is most reactive form of phosphorus and is highly polymerised.
- **38.** (c) Carbon has 2 electrons in their penultimate shell configuration so due to *d*-orbital in penultimate shell is false statement.
- **39.** (d) All other oxides of nitrogen except  $N_2O$  and NO are acidic in nature.
- **40.** (a)  $PbCl_4 < PbCl_2 < CaCl_2 < NaCl$  is the increasing order of ionic character.
- **41.** (a)  $Si + 3HCl \rightarrow SiHCl_3 + H_2$  silicon chloroform
- **42.** (a)  $KO_2$  is used in oxygen cylinder because it absorbs  $CO_2$  and increases  $O_2$  content. Super oxides reacts with water to give  $H_2O_2 \& O_2$ .
- **43.** (d)  $NaHCO_3$  and  $Na_2CO_3$  decomposes to give  $CO_2$  which extinguish fire.
- **44.** (d)  $CaF_2$  is insoluble in water.
- **45.** (a)  $PCl_3$  is most acidic.
- **46.** (b) CsCl > KCl > NaCl > LiCl
- **47.** (d)  $2Na_2S_2O_3 + I_2 \rightarrow Na_2S_4O_8 + 2NaI$ .
- **48.** (b) In general, higher the oxidation state, more is the covalent character of the oxide.  $I_2O_7 \otimes I_2O_5$  do not exist.  $I_2O_4$  is ionic in

- nature. It is infact iodyl iodate  $[IO]^+[IO_3]^-$ . The only covalent oxide is  $I_2O_5$ .
- **49.** (b)  $LiAlH_4$  act as reducing agent  $CH_3CHO + 2H \xrightarrow{LiAlH_4} CH_3CH_2OH$
- **50.** (c) When conc. HCl react with potassium chlorate to gives  $Cl_2 + ClO_2$ .
- **51.** (c)  $2H_2S + SO_2 \rightarrow 2H_2O + 3S$
- **52.** (b) Lithium nitrate on heating gives  $4LiNO_3 \xrightarrow{\Delta} 2Li_2O + 4NO_2 \uparrow +O_2 \uparrow$
- **53.** (d)  $CaCO_3 > NaHCO_3 > KHCO_3$  is the Increasing order of solubility.
- **54.** (c) Nitrolim is  $CaCN_2 + C$ .
- **55.** (a) The oxidation state of halogens is same +1. Therefore the acidic character depends only upon the electronegativity. Higher the electronegativity of the halogen more easily if will pull the electrons of O-H bond toward itself and hence, stronger is the acid.
- **56.** (c) Iridium [I, Z = 77] is not belong to noble gas.
- **57.** (c) Name and basicity of  $H_3PO_2$  is hypophosphorus acid & one.
- **58.** (c) In  $NO_2$  there are free electron so it is paramagnetic in nature.
- **59.** (a) Nessler's reagent is  $K_2HgI_4$ .
- **60.** (b) Helium was firstly discovered by willium Ramsay.
- **61.** (c) 1.66 Since inert gases are monoatomic.
- **62.** (c) In this reaction, phosphorus is simultaneously oxidised to  $NaH_2PO_2$  and reduced to  $PH_3$ . Hence this reaction is an example of disproportionation.

$$\stackrel{o}{P}_4 + 3NaOH + 3H_2O \rightarrow 3NaH_2\stackrel{+1}{P}O_2 + \stackrel{-3}{P}H_3$$

- **63.** (c) Reduction is accompanied by an increase in oxidation number of the reducing agent. C belong to IVA so the max-O.N. is +4. In  $CO_2$  the oxidation number of C is +4, which cannot be further increased. Hence,  $CO_2$  can not act as reducing agent.
- **64.** (b) When silicon tetra fluoride reacts with water  $H_2SiF_6$  and  $H_4SiO_6$  are formed

$$3SiF_4 \\ + 4H_2O \\ \rightarrow 2H_2SiF_6 \\ + \\ H_4SiO_4 \\ \text{White silicic acid}$$
 Silicon tetrafluoride Water

- **65.** (a)  $4O_3 + 6I_2(dry) \rightarrow 3I_4O_4$
- **66.** (c)  $Na(NH_4)HPO_4.4H_2O$  (Microcosmic salt)
- **67.** (b) Thermite mixture  $Fe_2O_3 + Al$

- **68.** (d) The colour of liquid  $O_2$  is pale blue.
- **69.** (a) Helium is not soluble in blood even under high pressure, a mixture of 80% helium and 20% oxygen is used instead of ordinary air by sea divers for respiration.
- **70.** (a)  $Mg_3N_2 + 6H_2O \rightarrow 3Mg(OH)_2 + 2NH_3$
- **71.** (a) Calcium cyanamide on treatment with steam produce  $CaCO_3 + NH_3$ .

#### Assertion & Reason

- 1. (b) Sulphate is estimated as  $BaSO_4$  because of high lattice energy in a group.
- **2.** (b) Fluorine is a strong oxidising agent than other halogens due to highest electronegativety..
- 3. (a) In  $HNO_3$  due to presence of two N-O bonds it is a stronger acid than  $HNO_2$ .
- **6.** (c) Bleaching action of chlorine carried by oxidation while bleaching action of  $SO_2$  carried by reduction.
- 7. (b) On heating its outermost electron transite to next energy level by which it become more reactive.
- **10.** (a) *K* and *Cs* emit electrons on exposure of light due to low ionisation potential.
- 11. (e) The lower value of bond dissociation energy of F-F bond due to longer inter electronic (electron electron) repulsion between the non-bonding electrons in the 2p orbitals of fluorine atom.
- 12. (a) It is fact that halogens are highly reactive as they have seven electrons in their outermost orbit and they want to stabilize by acquiring an electron. Therefor, they do not occur in free state. Here both assertion and reason are true and the reason is the correct explanation of assertion.
- 13. (a) Lithium forms lithium oxide. This is due to the fact that  $Li^+$  ion has smallest size and it has a strong positive field around it. Therefore, it stabilize  $O^{2-}$  ion with strong negative field around it. Thus, both assertion as well as reason are true.
- 14. (a) Liquid  $NH_3$  is used for refrigeration is true and it is due to the fact that is vaporises quickly and for vaporisation it takes up heat and cool the refrigerator. Hence assertion and reason both are true.
- **15.** (a) It is true that  $Al(OH)_3$ . The reason is that NaOH is a strong alkali, it dissolves  $Al(OH)_3$ . Which is amphoteric in nature and forms  $NaAlO_2$ .

$$Al(OH)_3 + NaOH \rightarrow NaAlO_2 + 2H_2O$$
Sodium meta aluminate

- **16.** (c) Boron is metalloid. Thus assertion is correct. Metalloids possess, metallic as well as nonmetallic nature. Hence, reason is false.
- 17. (b) It is correct that inert gases are monoatomic because for inert gases  $C_p/C_v=1.66$ .
- **18.** (c) When Mg is burnt in nitric oxide it continue to burn because during burning the heat evolved decompose NO to  $N_2$  and  $O_2$ . Oxygen thus, produced helps Mg to burn.

Here assertion is true but reason is false.

- 19. (d) Anhydrous  $BaO_2$  is not used for preparing  $H_2O_2$  because it reacts with  $H_2SO_4$  and the reaction ceases after some time due to formation of  $BaSO_4$  on  $BaO_2$ . Therefore, assertion and reason both are false.
- **20.** (d) Inorganic benzene, borazine is highly reactive while benzene is much less reactive.

Here, assertion is false, but reason is true.

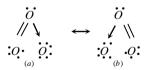
- 21. (a) The halogens absorb visible light due to which all halogens are coloured. Hence, both assertion and reason are true and reason is correct explanation.
- 22. (b) It is true that barium is not required for normal biological function in human beings and it is also true that it does not show variable oxidation state. It only shows +2 oxidation state.
- **23.** (d) The O-O bond length is shorter in  $O_2F_2$  than in  $H_2O_2$  due to higher electronegativity.  $H_2O_2$  is a non ionic compound. Here both assertion and reason are false.
- **24.** (d) Here both assertion and reason are false because  $PbI_4$  is not a stable compound and iodine can not stabilize higher oxidation states. Pb shows (II) oxidation state more frequently than Pb (IV) due to inert pair effect.
- 25. (b) Both assertion and reason are true but reason is not correct explanation of the assertion. Enamel the hardest substance of the body is composed of fluorine not magnesium. Magnesium is an essential element as it acts as a factor of many enzymes of glycolysis and a number of other metabolic reactions dependent upon ATP.
- **26.** (d) Both assertion and reason are false.

Radium is the rarest of all s-block elements comprising only  $10^{-10}$  percent of igneous rocks. Francium (s-block member) is radioactive; its long lived isotope  $^{223}$  Fr has a half life of only 21 minutes.

- **27.** (c) Assertion is true but reason is false. Due to high polarizing power of  $Li^+, LiCl$  is a covalent compound.
- **28.** (c) Assertion is true but reason is false. Be has fully filled  $2s^2$  orbital which gives a relatively more stable electronic configuration.
- **29.** (a) Both assertion and reason are true and reason is the correct explanation of assertion.
- **30.** (a) Both assertion and reason are true and reason is the correct explanation of assertion.
- **31.** (a) Both assertion and reason are true and reason is the correct explanation of assertion.
- 32. (c) Assertion is true but reason is false.

  Helium is a noble gas (Chemically inactive) but beryllium is a member of alkaline earth metals (Chemically active).
- 33. (b) Both assertion and reason are true but reason is not the correct explanation of assertion. Lattice energy of  $Na_2SO_4$  is less than its hydration energy but the lattice energy of  $BaSO_4$  exceeds its hydration energy.
- **34.** (a) Both assertion and reason are true and reason is the correct explanation of assertion.
- **35.** (a) Both assertion and reason are true and reason is the correct explanation of assertion.

  Presence of unpaired electrons in super oxides of alkali metals make them paramagnetic.
- **36.** (b) Both assertion and reason are true but reason is not the correct explanation of assertion. Nitrogen can not expand its octet due to the non availability of *d*-orbital.
- 37. (a) Both assertion and reason are true and reason is the correct explanation of assertion.Ozone is considered to be a resonance hybrid of the following two forms.



- 38. (b) Both assertion and reason are true but reason is not the correct explanation of assertion.
  Molecules of sulphuric acid are associated due to large number of intermolecular hydrogen bonding.
- **39.** (b) Both assertion and reason are true but reason is not the correct explanation of assertion.  $PCl_5$  is trigonal bipyramidal containing  $sp^3d$  hybridized P atom in liquid and gaseous state. Whereas in solid state it consists of tetrahedral  $PCl_4^+$  cation and octahedral  $PCl_6^-$  anion.

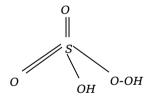
- **40.** (b) Both assertion and reason are true but reason is not the correct explanation of assertion. In case of  $NI_3$ , the lone pair moment adds on the resultant of the N-I moments but in case of  $NF_3$ , the lone pair moment on partly cancels the resultant N-F moments.
- 41. (a) Both assertion and reason are true and reason is the correct explanation of assertion.

  The ignition temperature of white phosphorus is low (About 30°C). In air it readily catches fire giving dense fumes of phosphorus pentoxide. It is therefore, kept in water.
- 42. (c) Assertion is true but reason is false.B does not have vacant *d*-orbitals as second shell is the outermost shell.
- **43.** (a) Both assertion and reason are true and reason is the correct explanation of assertion.
- **44.** (d) Both assertion and reason are false. Si-Si bonds are weaker than Si-O bonds Si has no tendency to form double bonds with itself.
- **45.** (c) Assertion is true but reason is false. S atoms in  $S_8$  molecule undergo sp hybridization and contain two lone pairs of

hybridization and contain two lone pairs of electrons on each and exists as staggered 8 atom rings.

**46.** (a) Both assertion and reason are true and reason is the correct explanation of assertion.

This can be explained through structure of Caro's acid (Peroxomonosulphuric acid).



Oxidation no. of S = x, oxidation no. of H = +1,

Oxidation no. of oxygen in peroxo linkage = -1,

Oxidation no. of other oxygen atoms = -2 (each) 2+x-6-2=0 or x=+6.

- 47. (d) Both assertion and reason are false.

  The melting point / boiling point of noble gases are quite low. The inter particle forces among noble gases are weak Vander Waal's forces.
- **48.** (b) If both assertion and reason are true and reason is the correct explanation of assertion. S atom in both  $SO_2$  and  $SO_3$  is  $sp^2$  hybridized but it contains a lone pair of electrons in  $SO_2$ .
- 49. (d) Both assertion and reason are false.

Calcium carbide on hydrolysis gives acetylene. Calcium carbide contains  $C_2^{2-}$  anion.

- **50.** (a) Both assertion and reason are true and reason is the correct explanation of assertion.
- **51.** (e) Assertion is false but reason is true.

  In lab, hydrogen is generally prepared by the reaction of zinc with dilute hydrochloric acid.
- **52.** (b) Both assertion and reason are true but reason is not the correct explanation of assertion. The relatively inert behaviour of diprotium at room temperature is due to the high enthalpy of H-H bond, being the highest for a single bond between any two elements.
- 53. (d) Both assertion and reason are false.

  Water can be easily transformed from liquid solid and to gaseous states. The distribution of water over the earth's surface is not uniform. The desert region have no permanent surface water while the oceans cover vast areas.
- 54. (b) Both assertion and reason are true but reason is not the correct explanation of assertion.
  The structure of ice is open due to hydrogen bonding which makes ice less dence than liquid water at the same temperature.
- 55. (c) Assertion is true but reason is false.

  The water molecules are joined together in an extensive three dimensional network in which each oxygen atom is bonded to four hydrogen atoms two by hydrogen bonds and two by normal covalent bonds in a near tetrahedral configuration. This situation does not exist for molecules like *NH*<sub>3</sub> and *HF*.
- 56. (a) Both assertion and reason are true and reason is the correct explanation of assertion.Hard water contain soluble calcium and magnesium salt like bicarbonates, chlorides and sulphates.
- 57. (a) Both assertion and reason are true and reason is the correct explanation of assertion.

  To stop decomposition  $H_2O_2$  is stored in wax-lined glass or plastic vessels in the presence of stabilizers like urea.

58. (a) Both assertion and reason are true and reason

- is the correct explanation of assertion. At any given instant in liquid water at room temperature, each water molecule forms hydrogen bonds with an average 3.4 other water molecules. The  $H_2O$  molecules are in continuous motion so hydrogen bonds are constantly and rapidly broken and formed. In ice  $H_2O$  molecules are, however fixed in the
- 59. (d) Both assertion and reason are false.

space lattice.

Calgon is used for making  $Ca^{2+}$  and  $Mg^{2+}$  ions present in hard water ineffective. It forms soluble complex with  $Ca^{2+}$  and  $Mg^{2+}$  ions.

**60.** (b)  $SO_2$  shows both oxidising as well as reducing nature.

The reaction given in assertion is due to oxidising nature of  $SO_2$ .

$$2H_2S + SO_2 \rightarrow 2H_2O + 3S$$

- **61.** (a) (i) Due to smaller size of F; steric repulsions will be less in  $SiF_6^{2-}$ .
  - (ii) Interaction of F lone pair electrons with Si is stronger than that of chlorine lone pairs.
- **62.** (b) Borax bead test is not suitable for Al(III) because its oxidising as well as reducing flame is colourless in both hot as well as cold. Alumina is insoluble in water as they exist in hydrated form like  $Al_2O_3.2H_2O$ ,  $Al_2O_3.H_2O$  etc.
- **63.** (c)  $SeCl_4$  possess see saw geometry, which can be regarded as a distorted trigonal bipyramidal structure laving one lone pair (lp) of electrons in the basal position of the trigonal bipyramidal. See-saw geometry of  $SeCl_4$  molecules arises due to the  $sp^3d$  hybridisation of the central atom. The distortion in shape is due to the presence of one lone pair of electrons.
- **64.** (b) Due to the ease with which it can liberate nascent oxygen,  $O_3$  acts as a powerful oxidising agent.

