

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) KHCO₂ is less soluble than NaHCO₂
- 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 13.
 - (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) CO₂
- (b) Water vapour
- (c) CO
- (d) No gas
- Chile saltpetre is
- [DPMT 1984; CPMT 1986, 89; CET Pune 1998; MP PMT 2003]
 - (a) $NaNO_3$
- (b) Na_2SO_4
- (c) KNO_3
- (d) Na_2SO_3
- A mixture of KCl and KF is added to sodium 16. 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

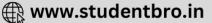
Sodium metal can be stored under

[CPMT 1972, 85; BHU 1983]

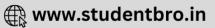
- (a) Benzene
- (b) Kerosene
- (c) Alcohol
- (d) Toluene







19.	_	method of preparing		The colour of hydrogen	is [IIT 1980]
		he action of <i>HCl</i> and[JIPME	R 2000]	(a) Black	(b) Yellow
	(a) Al	(b) <i>K</i>		(c) Orange	(d) None of these
	(c) Fe	(d) Zn	32.		owing salts gives aqueous
20.		y and other considerations ving alkali metal chlorides		solution which is weakl	=
		highest melting point[AIEEE		(a) NaHCO ₃	(b) NaHSO 4
	(a) LiCl	(b) NaCl		(c) NaCl	(d) NH_4HCO_3
	(c) KCl	(d) RbCl	33.	An example for a doubl	
21.	The correct formula of l			(a) Silver nitrate	(b) Mohr's salt
		(b) Na_2SO_4			ide (d)Cupromonium sulphate
	(c) $Na_2S_2O_3.4H_2O$		34.	The elements of group flame of Bunson burner	IA provide a colour to the due to [AIIMS 1987]
22.		nly used to determine		(a) Low ionization pote	
	_	netrically is [AIIMS 2003]		(b) Low melting point	
	(a) Oxalic acid			(c) Softness	
	(b) Disodium salt of ED	TA		(d) Presence of one e	lectron in the outermost
	(c) Sodium citrate		orbit	t	
	(d) Sodium thiosulphate	2	35.	Which of the following	is the smallest cation
23.	K_2CS_3 can be called pot	tassium [CPMT 1972, 74]			[MP PMT 1993]
	(a) Thiocyanate	(b) Thiocarbonate		(a) <i>Na</i> ⁺	(b) Mg^{+2}
	(c) Thiocarbide	(d) Sulphocyanide		(c) Ca ⁺²	(d) Al^{+3}
24.	Which is most basic in o		36.		may be arranged in the
	(a) <i>RbOH</i> (c) <i>NaOH</i>	(b) <i>KOH</i> (d) <i>LiOH</i>		_	their standard electrode
25.	When washing soda is h			potentials as (a) <i>K</i> , <i>Ca</i> , <i>Li</i>	[CPMT 1990] (b) <i>Li</i> , <i>K</i> , <i>Ca</i>
25.	(a) CO is released	[AFMC 2005]			
	(b) $CO + CO_2$ is released	d		(c) Li, Ca, K	(d) Ca, Li, K
	(c) CO_2 is released		37•	Alkali metals lose electr (a) s-orbitals	ons in [CBSE PMT 1990] (b) <i>p</i> -orbitals
	(d) Water vapour is rele	eased		(c) <i>d</i> -orbitals	(d) f-orbitals
26.	Which of the following i		38.		acts with nitrogen directly
	(a) All carbonates are se		3		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	er			(c) K	(d) <i>Rb</i>
	(c) Carbonates of <i>Ca</i> , <i>S</i>	<i>Sr</i> , <i>Ba</i> are soluble in water	39.	-	has density greater than
	(d) All carbonates are in	nsoluble		water	[MP PET 1994]
27.	Nitre is	[CPMT 1986]		(a) Li	(b) Na
	(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 t	he preparation of		water, is made use of	[MP PMT 1994]
		[CPMT 1985]		(a) In drying of alcohol:	
	(a) Slaked lime	(b) Baryta		(b) In drying of benzene	
	(c) Sodium	(d) Caustic soda		(c) In drying of ammon	
29.	Potash alum is a	[CPMT 1986; MNR 1981]		(d) As a general drying	•
	(a) Complex salt	(b) Acid salt	41.	Which of the following	has smaller size [RPET 2003]
20	(c) Double salt	(d) Normal salt ustrial manufacturing of		(a) H	(b) He ⁺
30.	sodium carbonate is known	own as [CPMT 1978, 86; MP P	PMT 1995	(a) H	
	(a) Castner process	(b) Haber's process	223	-(c) ₁ H	(d) Li^{2+}
	(c) Le-blanc process	(d) Chamber process			



- 42. combines with HF to form KHF_2 . The compound contains the species [IIT 1996]
 - (a) K^+ , F^- and H^+
- (b) K^+ , F^- and HF
- (c) K^+ and $[HF_2]^-$
- (d) $[KHF]^+$ and F^-
- Which alkali metal is most metallic in character

[MH CET 2001]

(a) K

- (b) Cs
- (c) Na
- (d) Li
- The property of hydrogen which distinguishes it from other alkali metals is [MP PET 1996]
 - (a) Its electropositive character
 - (b) Its affinity for non-metals
 - (c) Its reducing character
 - (d) Its non-metallic character
- Which of the following reacts with water with 45. high rate

[AFMC 1995]

- (a) *Li*
- (b) K
- (c) Na
- (d) Rb
- The valence shell electronic configuration of alkali metals is

[MP PET 1996; UPSEAT 2001]

- (a) ns^2np^1
- (b) ns¹
- (c) $(n-1)p^6ns^2$
- (d) $(n-1)d^2ns^2$
- Alkali metals are
- [MP PMT 1996]
- (a) Li, Na, Be, Mg, Cs
- (b) Li, Na, K, Rb, Cs
- (c) Na, K, Mq, Ca, Rb
- (d) K, Rb, Cs, Ba, Sr
- 48. The atomic number of an element is 11. Its oxide will be

[MP PMT 1996]

- (a) Acidic
- (b) Basic
- (c) Acid and basic both (d) Neutral
- The commercial production of sodium carbonate is done by

[CPMT 1982; MP PMT 1996]

- (a) Lead-chamber process
- (b) Haber's process
- (c) Solvay's process
- (d) Castner's process
- 50. Alkali metals are strong reducing because
 - (a) These are monovalent
 - (b) Their ionisation potential are very high
 - (c) Their standard electrode potential are very much negative
 - (d) These are metals
- Which of the following statement about LiCl and 51. NaCl is correct [Kurukshetra CEE 2002]

- (a) LiCl has higher melting point than NaCl
- (b) LiCl dissolves in water whereas NaCl does not
- (c) LiCl would ionize in water more than NaCl
- (d) Fused LiCl would be less conducting than fused NaCl
- In the Castner's process for the extraction of sodium, the anode is made of......metal. [EAMCET 2003]
 - (a) Copper
- (b) Iron
- (c) Sodium
- (d) Nickel
- 53. Which of the following s-block elements forms nitride

[RPET 2003]

[Pb. PMT 2001]

- (a) Ba
- (b) Be
- (c) Ca
- (d) Li
- Tincal is 54.
- (b) NaNO 3
- (c) NaCl

(a) $Na_2CO_3.10H_2O$

- (d) $Na_2B_4O_7.10H_2O$
- Which has minimum solubility [BHU 2003] 55.
 - (a) Br_2S_3
- (b) Ag_2S
- (c) CoS
- (d) PbS
- **56.** Cryolite helps in
- [BHU 2003]
- (a) Lowering the melting point
 - (b) Increasing the melting point
 - (c) Increasing the electrical conductivity
 - (d) Decreasing the electrical conductivity
- In certain matters lithium differs from other alkali metals, the main reason for this is[MP PET/PMT 199
 - (a) Small size of Li atom and Li^+ ion
 - (b) Extremely high electropositivity of Li
 - (c) Greater hardness of Li
 - (d) Hydration of Li^+ ion
- Acidified potassium permanganate solution is 58. decolourised by
 - (a) Bleaching powder
- (b) Microcosmic salt
- (c) Mohr salt
- (d) White vitriol
- Which one of the following is used as a disinfectant in water treatment [NDA 1999]
 - (a) Alum
- (b) Charcoal
- (c) Kieselguhr
- (d) Potassium

permanganate

60. Sodium thiosulphate is used in photography

[UPSEAT 1999]

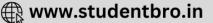
- (a) To convert metallic silver into silver salt
- (b) AgBr grain is reduced to non-metallic silver
- (c) To remove reduced silver
- (d) To remove undecomposed AqBr in the form of $Na_3[Ag[S_2O_3)_2]$ (a complex salt)
- Composition of borax is

[UPSEAT 2001;04]

- (a) $Na_{2}B_{4}O_{7}.4H_{2}O$
- (b) $Na_2B_4O_7.10H_2O$





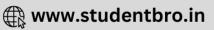


758 s and p-Block Elements (c) $NaBO_{2}$ (d) Na_2BO_3 (b) Does not show regular variation (c) Increases as we go down the group 62. When sodium dicarbonate is heated strongly for calcined in a kiln, it forms[CPMT 2000; KCET (Med.) 2000] (d) Decreases as we go down the group (b) Na_2CO_3 (a) Na Which of the following properties is not true for an alkali metal [Pune CET 1998] (c) NaCO₃ (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 The word 'alkali' is used for alkali metals (d) Low electronegativity 64. indicates Which of the following alkali metals has the 73. tendency for the half [RPMT 1999] biggest reaction, (a) Ash of the plants (b) Metallic nature $M(g) \rightarrow M^+(aq) + e^-$ [DPMT 2001] (c) Silvery lusture (d) Active metal (a) Lithium (b) Sodium 65. Potassium nitrate is called [RPMT 1999] (c) Cesium (d) Potassium (a) Mohr's salt (b) Gypsum Which one of the following metalic hydroxides (d) Chile salt petre (c) Indian 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 (d) $Pb(OH)_{2}$ (c) $Fe(OH)_3$ Solvay process [Roorkee 1999] Which one of the following on heating will not give CO_2 (a) NaCl, CO and NH_3 [NDA 1999; BHU 2000] (b) $NaCl, CO_2$ and NH_3 (a) CaCO₃ (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 ° 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^{Θ} (b) NaOH (a) Na_2O (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 melts resolidifies and then liberates a gas. It may be[DPMT 2002] (a) Li (b) Na (d) Cs (c) K (a) MnO_2 (b) Al_2O_3 With the increase in atomic weights, melting (c) $KMnO_4$ (d) KClO₃ 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_3 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] (d) Irreversible and endothermic

(a) Increases at first and then decreases

				s and p-block Lientents /59
81.	_	ocyanide crystals are heated		(c) CH ₃ COONa (d) None of these
	is	phuric acid, the gas evolved	91.	A fire of lithium, sodium and potassium can be extinguished by [DCE 2003]
		SE PMT PMT 1999; KCET 2000]		(a) H_2O (b) Nitrogen
	(a) Ammonia	(b) Sulphur dioxide		
_	(c) Carbon dioxide	(d) Carbon monoxide		(c) CO_2 (d) Asbestos blanket
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	ts		(a) Na (b) Mg
	(c) Low oxidation potential	entials		(c) Al (d) Si
	(d) High ionization po	tentials	93.	Aluminium reacts with caustic soda to form [DCE 200
33.		compound of an element of		(a) Aluminium hydroxide
		te X gives a violet colour in		(b) Aluminium oxide
	flame test, <i>X</i> is	0- 06 CDMT0- DCT1		(c) Sodium meta-aluminate
		85, 86; CPMT 1985; DCE 2000]		(d) Sodium tetra aluminate
	(a) LiCl	(b) NaCl	94.	Alkaline earth metals are denser than alkali metals,
0.4	(c) KCl	(d) None		because metallic bonding in alkaline earth's metal,
84.		ring alkali metal ions has In aqueous solution [KCET 2000	1	is [BHU 2004]
	(a) Rb^+	(b) Cs ⁺	,	(a) Stronger (b) Weaker
		` '		(c) Volatile (d) Not present
	(c) Li ⁺	(d) Na ⁺	95.	Which of the following is a false statement [CPMT 26
35.	chemical behaviour be	arities to magnesium in its		(a) Fluorine is more electronegative than chlorine
		ecause [Pb. PMT 2000] eater electronegativity and		(b) Nitrogen has greater IE_1 than oxygen
	similar polarizing			(c) Lithium is amphoteric
		electronegativity and lower		(d) Chlorine is an oxidising agent
	polarizing power	creeti onegativity and lower	96.	Which is most basic in character [UPSEAT 2004]
	(c) Similar size, same	electronegativity and		(a) CsOH (b) KOH
	similar high polari			(c) NaOH (d) LiOH
	(d) None of these		97.	Photoelectric effect is maximum in [AFMC 2004]
36.		following is the most	٠,٠	(a) Cs (b) Na
	electropositive elemen	it [Pb. PMT 2000]		(c) K (d) Li
	(a) Calcium	(b) Chlorine	۵R	A metal M reacts with N_2 to give a compound ' A '
	(c) Potassium	(d) Carbon	30.	(M_3N) . 'A' on heating at high temperature gives
37.	-	n sodium chloride leads to		back 'M' and 'A' on reacting with H_2O gives a gas
	the formation of	[KCET 1990]		'B'. 'B' turns $CuSO_4$ solution blue on passing
	(a) Na and H_2	(b) Na and O_2		
	(c) H_2 and O_2	(d) Na and Cl_2		through it. A and B can be [DCE 2003]
88.	When sodium bicarbo	onate is heated the product		(a) Al and NH_3 (b) Li and NH_3
	obtained is	1		(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 ₃	(d) $Na_2(HCO_3)$		forms 'Y'. On passing an excess of ${\it CO}_2$ through 'Y'
39.	5	g is a use of alum [CPMT 2004]		in water, a clear solution, ${}^{\prime}Z^{\prime}$ is obtained. On
9.				boiling 'Z', compound 'X' is reformed. The
	(a) Making explosives(c) Water softening	(b) Bleaching clothes(d) All of the above		compound 'X' is [CBSE PMT 2004]
	•			(a) Na_2CO_3 (b) K_2CO_3
90.	hydrolysed in water	owing salt does not get		(c) $Ca(HCO_3)_2$ (d) $CaCO_3$
	ing at ony oca in water	[CPMT 2004]	100.	mongst $LiCl$, $RbCl$, $BeCl_2$ and $MgCl_2$ the compounds
	(a) KClO ₄	(b) NH_4Cl		with the greatest and least ionic character respectively are





	700 s and p-Bloc	k Elements			
		[Pb. CET 2004]	111.		ved in water, the sodium ion
	(a) LiCl and RbCl	(b) $MgCl_2$ and $BeCl_2$		is	[CDMT 1090]
	(c) RbCl and BeCl ₂	(d) $RbCl$ and $MgCl_2$		(a) Oxidised	[CPMT 1989] (b) Reduced
101.	Salt cake is			(c) Hydrolysed	(d) Hydrated
	(a) Sodium sulphate		112	Sodium metal cannot	
	(b) Sodium chloride(c) Sodium bisulphite		112.	Soutum metal calmot	[CPMT 1985, 88, 94]
	(d) Sodium sulphate an	nd Sodium chloride		(a) Benzene	(b) Kerosene
102	Globar salt is	[BHU 1983; CPMT 1988, 91;		(c) Alcohol	(d) Toluene
102.	Global balt is	IIT 1985; MP PET 2000]	112		s is used for the preparation
	(a) $MgSO_4.7H_2O$	(b) $CuSO_4.5H_2O$	113.	of	o to used for the preparation
	(c) $FeSO_4.7H_2O$	(d) $Na_2SO_4.10H_2O$			[CPMT 1985; BHU 1986]
103.	· -	flame by sodium salts is		(a) Caustic soda	(b) Caustic potash
3.	9	[CPMT 1980; MP PET 1986]		(c) Baryta	(d) Slaked lime
	(a) Light red	(b) Golden yellow	114.	When CO is passed	over solid NaOH heated to
	(c) Green	(d) Pink		$200^{o}C$, it forms	[MP PMT 1985]
104.	Solvay's process is used			(a) Na_2CO_3	(b) NaHCO ₃
		[CPMT 1982; AIIMS 1987]		(c) HCOONa	(d) None
	(a) Ammonia	(b) Sodium bicarbonate	115.		red by electrolysis of brine
	(c) Sodium carbonate	(d) Calcium carbonate			s of the reaction are[KCET 1990]
105.	gives	n a current of dry ammonia		(a) Cl_2 and H_2	(b) Cl_2 and $Na - Hg$
	gives	[NCERT 1981; KCET 2000]		(c) Cl_2 and Na	(d) Cl_2 and O_2
	(a) Sodium nitrite	(b) Sodium hydride	116	_	s manufactured by Solvay
	(c) Sodium amide	(d) Sodium azide	110.		that are recycled are[KCET 1993; DO
106.	Washing soda is			(a) CO_2 and NH_3	(b) CO_2 and NH_4Cl
	[CPMT 1982;	DPMT 1982; CBSE PMT 1990;		(c) NaCl, CaO	
		MP PMT 1987, 96]			(d) CaCl ₂ , CaO
	(a) $Na_2CO_3.10H_2O$	(b) $Na_2CO_3.H_2O$	117.		icts, obtained in the Solvay ring sodium carbonate, are[KCET 19
	(c) $Na_2CO_3.5H_2O$	(d) Na_2CO_3		(a) Quick lime and CO	_
107.		decolourise and purify oils		•	-
	is			(b) $NaHCO_3$ and NH_4	Cl
	(a) Codium combonato	[MP PMT 1987] (b) Sodium chloride		(c) NH_4Cl solution ar	nd quick lime
	(a) Sodium carbonate(c) Sodium hydroxide	(d) Sodium sulphate		(d) $NaHCO_3$ and CO_2	
108	•	n sea water is[MP PMT 1998]	118.	In the preparation of	sodium carbonate, which of
100.	(a) $MgCl_2$	(b) <i>NaCl</i>		the following is used	[AFMC 1992]
				(a) Slaked lime	(b) Quick lime
	(c) $MgSO_4$	(d) CaSO ₄		(c) Lime stone	(d) NaOH
109.	The metallic lustre explained by	exhibited by sodium is	119.	_	s are left in open air, they round each crystal as[CPMT 1974]
	(a) Diffusion of andium	[IIT 1987]		(a) They start melting	
	(a) Diffusion of sodium			(b) They absorb moist	ture from air
	(b) Oscillation of loose			•	r to form a liquid compound
	(c) Excitation of free p			(d) They absorb CO_2 :	
	(d) Existence of body c		120	_	
110.	temperature is	acts with water at room	120.		acts with SO_2 in aqueous
	=	MP PMT 1996; MP PET 1998]		medium to give	[MD DMT 1082 8=1
	(a) Copper	(b) Iron		(a) NaHSO 3	[MP PMT 1982, 85]
	(c) Magnesium	(d) Sodium		_	(b) Na_2SO_3
	(c) magnesiam	(a) ooaram		(c) $NaHSO_4$	(d) Na_2SO_4

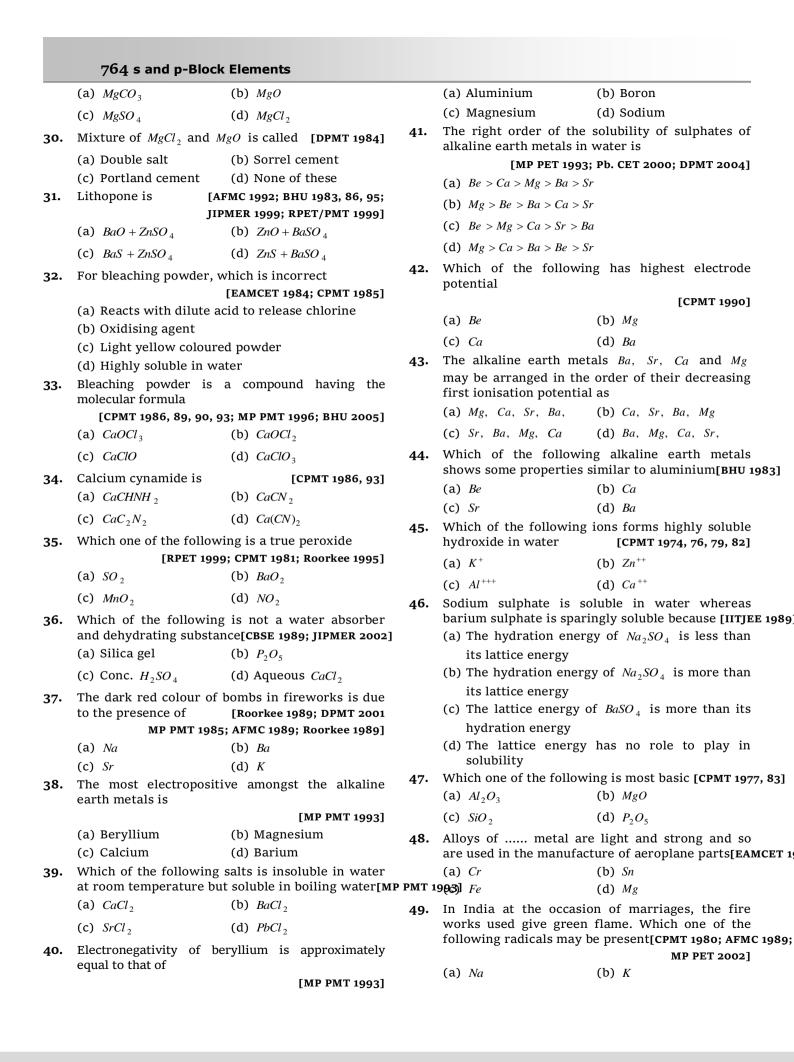
	Paleta and 1 1	F		(1-) 100/ 111 6	77
121.	Baking soda is	[CPMT 1974, 78, 79, 91;		(b) 10% solution of KO	
	CPMT 1973; RPET 1999; AFM	al MEE 1995; AIIMS 1996;		(c) 10% solution of <i>Ca</i> (
) NaHCO ₃		(d) 10% solution of Na_2	$_{2}CO_{3}$
		K_2CO_3	131.		colour to Bunsen flame
400				because of	[RPMT 1997]
122.	Soda ash is	[KCET 1993]		(a) Low ionisation pote	
	2 3 2) NaOH		(b) Sensitivity	
	(c) Na_2CO_3 (d)	NaHCO ₃		(c) Sublimation	
123.	Soda lime is	[KCET 1993]		(d) Absorbed high radia	ition
	(a) NaOH (b) CaO	132.		ess of <i>NaOH</i> solution, the
	(c) NaOH and CaO (d	Na_2CO_3		compound obtained is	[RPMT 1997]
124.	Molten sodium is used in n	uclear reactors to[KCET 19	89]	(a) $Sn(OH)_2$	(b) Na_2SnO_3
	(a) Absorb neutrons in ord	ler to control the chain		(c) Na_2SnO_2	(d) SnO_2
react			133.	Identify the correct stat	
	(b) Slow down the fast neu				can be prepared and ysing an aqueous solution
	(c) Absorb the heat genera	-		of sodium chloride	ysing an aqueous solution
	(d) Extract radio-isotopes			(b) Elemental sodium is	s a strong oxidizing agent
125.	Squashes are stored by add	_		(c) Elemental sodium is	s insoluble in ammonia
) KCl		(d) Elemental sodium is	s easily oxidized
mate	2 3) Sodium	134.	Calcium is obtained by	[CBSE PMT 1997]
	bisulphite	C O 5H O) is used in		(a) Roasting of limeston	
126.	Sodium thiosulphate (Na_2 photography to	$S_2O_3.5H_2O_1$ is used in [CPMT 1972, 74, 79;		(b) Electrolysis of solution H_2O	tion of calcium chloride in
		nar CEE 1995; MNR 1995]		(c) Reduction of calciur	n chlorido with carbon
	(a) Reduce silver bromide				olten anhydrous calcium
	(b) Convert metallic silver		chlo	-	onen annyarous calerain
	(c) Remove undecompose		135.		e solution is electrolysed,
	silver thiosulphate com			=	at the cathode is[Kurukshetra CEI
	(d) Remove unreduced silv	er		(a) Oxygen	(b) Hydrogen
127.	Which of the following pair	can't exist in solution	106	(c) Chlorine	(d) Air of fused sodium chloride,
		[IIT 1986; DCE 1999]	130.	the anodic reaction is	[KCET 1998]
	(a) $NaHCO_3$ and $NaOH$ (b)) Na_2CO_3 and $NaOH$		(a) Reduction of sodium	
	(c) Na_2CO_3 and $NaCl$ (d) NaHCO ₃ and NaCl		(b) Oxidation of sodium	ions
128.	Sodium thiosulphate is pre-	pared by [IIT 1996]		(c) Reduction of chlorid	le ions
	(a) Reducing Na_2SO_4 solut	•		(d) Oxidation of chlorid	
	(b) Boiling Na_2SO_3 solution	-	137.		does not participate in the
medi	·	on with 5 in aranic			manufacture of Na_2CO_3 [EAMCET
	(c) Neutralising $H_2S_2O_3$ so	olution with NaOH		(a) NH_3	(b) NaCl solution
	(d) Boiling Na_2SO_3 solution			(c) CO_2	(d) H_2SO_4
medi	ium			The colour of the precing NaOH solution to HgCl	pitate produced by adding [KCET 1998]
129.	When <i>NaOH</i> is prepared, t	_	96]	(a) Yellow	(b) Black
		H_2		(c) Brown	(d) White
	(c) O_2 (d)	$) H_2O$	139.		ectrolysis of fused NaCl is
130.	What is lye	[BHU 1997]			IC 1999; Kerala (Mea.) 2002]
	(a) 10% solution of NaCl			(a) Down's cell	(b) Castner cell



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; (c) Potash alum (d) Ammonium 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. (c) Cyanide process (d) Both (b) and (c) [EAMCET 1978; CPMT 1988; CET Pune 1998; 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. Which of the following ore contains both 5. (c) Diabetes (d) Anaemia magnesium and calcium [MDAT Bihar 1984; MP PET 2003] 145. Ferric alum the composition (a) Magnesite (b) Dolomite [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) $CaSO_4.2H_2O$ (a) Na_2CO_3 (b) Na_2O_2 (b) $BaSO_4.2H_2O$ (c) Na_2O (d) Both (b) and (c) (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) Cl₂ (c) Dehydration 148. Pyrolusite is [DPMT 2002] (d) Hydration to yield another hydrate (b) Sulphur ore (a) Carbonate 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 (c) Metal is taken out by spoons (a) Improve the electrical conduction (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 Alkaline earth metals (c) K_2CO_3 (d) Rb_2CO_3

11.	The outer electronic	configuration of alkaline			[NCERT 1982]
	earth metal is			(a) Dissociation of b	peryllium carbide
	[BHU 1980;	CPMT 1985, 93; MP PAT 1993]		(b) Electrolysis of fu	used beryllium chloride
	(a) ns^2	(b) ns^1		(c) Reduction of ber	ryllium oxide with carbon
	(c) np 6	(d) nd^{10}		(d) Reduction of	beryllium halide with
12.	Metallic magnesium is	prepared by	mag	BACⁱ 197 3, 77]	
	(a) Reduction of MgO	by coke	21.	Mark the incorrect s	statement
	(b) Electrolysis of aqu	eous solution of $Mg(NO_3)$		(a) Lithopone is che	eap and possess good covering
	(c) Displacement of	Mg by iron from $MgSO_4$	pow		
solu				(b) Lithopone is yell	
	(d) Electrolysis of mol	ten MgCl ₂		(c) Lithopone is p sulphide and zin	prepared by mixing bariun Ic sulphate
13.		, Ca and Sr of group II A.		=	nixture of barium sulphate and
		ne least ionic chloride would		zinc sulphide	-
	be formed by	INCERT 1000, CRWT 1000	22.	Pure anhydrous Mg	Cl_2 can be prepared from the
	(a) <i>Be</i>	[NCERT 1980; CPMT 1980] (b) Mg		hydrated salt by	[CPMT 1986; MP PMT 1989
	(c) Ca	_		(a) Heating the hydr	rate with coke
1.4	Which one of the follow	(d) Sr		(b) Heating the hydr	rate with Mg ribbon
14.	(a) CaF_2	(b) CaO		(c) Melting the hydr	rate
				(d) Heating the h	ydrate to red heat in ar
	(c) H_2F_2	(d) $CaCO_3$		atmosphere of <i>H</i>	
15.	Which one is known as	· ·	23.		obtained by the interaction o
	(a) $BaSO_4$	(b) $BaCl_2.2H_2O$			T 1972, 78, 89; 2002; DPMT 1983
	(c) <i>BaO</i>	(d) BaCO ₃		(a) Conc. solution of	$f Ca(OH)_2$
16.	Which of the following	g sulphates have the highest		(b) Dilute solution of	of $Ca(OH)_2$
	solubility in water[EAR	MCET 1980,84,85; MP PMT 1994;		(c) Dry calcium oxid	de
	Kurukshetra CEE 19	98; AFMC 1990; MP PET 1994]		(d) Dry slaked lime	
	(a) $MgSO_4$	(b) $BaSO_4$	24.	Deep pink colour is	given to flame by the salts of
	(c) CaSO ₄	(d) $BeSO_4$		(a) Strontium	(b) Potassium
17.	The composition form	ulae of gypsum is		(c) Zinc	(d) Barium
	-	, 78, 82; DPMT 1982; IIT 1978;	25.		which colour when put in a
	MNR 19	81; MP PMT 1996; RPMT 1997]		flame (a) Brick red	(b) Green
	(a) $(CaSO_4)_2.H_2O$	(b) $2CaSO_4$		(c) White	(d) Pink
	(c) $CaSO_4.2H_2O$	(d) $2CaSO_4.H_2O$	26.		ed from the following ore
18.	Mortar is a mixture of	[EAMCET 1998; AIIMS 2000]	_0.	Thoopinio to obtain	[Roorkee 1995]
	(a) CaCO ₃ , sand and v			(a) Calcium superpl	nosphite (b)Calcium phosphid
	(b) Slaked lime and wa			(c) Potassium phosp	ohide (d) Calcium
	(c) Slaked lime, sand a			hypophosphide	
	(d) $CaCO_3$ and CaO	and water	27.	Calcium is obtained	_
	_				IT 1980; CPMT 1996;AIIMS 2001
19.		on heating to about 120°C		(a) Roasting of lime	
	forms a compound	which has the chemical	MCET	(b) Reduction of Car	Cl ₂ with carbon
	composition represent	ed by[CPMT 1978, 82, 88, 90; EADPMT 1982, 83; NCERT 1979]	MICEI	'(ć)'Electrolysis of a	solution of $CaCl_2$ in water
	(a) CaSO ₄	(b) $2CaSO_4.H_2O$		(d) Electrolysis of m	nolten CaCl ₂
	(c) $CaSO_4.H_2O$	(d) $2CaSO_4.3H_2O$	28.	Which element poss	esses biggest atomic radii
	· -			(a) <i>P</i>	(b) <i>Si</i>
20.	The highly efficient beryllium is	nt method of obtaining		(c) Al	(d) <i>Mg</i>
	J		29.	Magnesia is	







s and p-Block Elements 765 (d) Be, Mq, Ca, K, Rb

 $CaCO_3 \square CaO + CO_2$ reaction in a line goes to completion because [AFMC 2005]

(d) Ca

(a) CaO does not react to CO_2 to give $CaCO_3$

- (b) Backward reaction is very slow
- (c) CO_2 formed escapes out
- (d) None of these

The wire of flash bulb is made of [CPMT 1988]

(a) Mg

(c) Ba

50.

- (b) Cu
- (c) *Ba*
- (d) Ag

52. Bone ash contains

[KCET 1992]

- (a) *CaO*
- (b) CaSO 4
- (c) $Ca_3(PO_4)_2$
- (d) $Ca(H_2PO_4)_2$

53. A substance absorbs CO_2 and violently reacts with water. That substance is

- (a) CaCO₃
- (b) CaO
- (c) H_2SO_4
- (d) ZnO

54. Setting of cement is an

[DPMT 1984]

- (a) Exothermic reaction
 - (b) Endothermic reaction
 - (c) Neither exothermic nor endothermic
 - (d) None of these
- 55. Which is quick lime

[EAMCET 1993]

- (a) $Ca(OH)_2$
- (b) CaO
- (c) CaCO₃
- (d) $Ca(OH)_2 + H_2O$

56. A major constituent of portland cement except lime is

[CPMT 1982]

- (a) Silica
- (b) Alumina
- (c) Iron oxide
- (d) Magnesia

Portland cement is manufactured by using [CPMT 1986]

- (a) Lime stone, clay and sand
- (b) Lime stone, gypsum and sand
- (c) Lime stone, gypsum and alumina
- (d) Lime stone, clay and gypsum

58. Identify the correct statement [CBSE PMT 1995]

- (a) Gypsum contains a lower percentage of plaster of calcium than plaster of paris
- (b) Gypsum is obtained by heating plaster of paris
- (c) Plaster of paris can be obtained by hydration of gypsum
 - (d) Plaster of paris is obtained by partial oxidation of gypsum
- 59. Which of the following decreases on going gradually from Be to Ba (in periodic table)
 - (a) Basic character of hydroxides
 - (b) Solubility of sulphates in water
 - (c) Solubility of hydroxides in water
 - (d) Strength of elements as reducing agent
- 60. Alkaline earth metals are [MP PMT 1996]

(a) Li, Be, K, Mq, Ca

- (b) Be, Mq, Ca, Sr, Ba
- (c) Be, K, Mg, Ca, Sr

Which of the following substances is used in the laboratory for fast drying of neutral gases[AIIMS 1998; AF

- (a) Sodium phosphate
- (b) Phosphorus pentoxide
- (c) Sodium sulphate
- (d) Anhydrous calcium chloride
- Which of the following can be represented by the configuration $[Kr]5s^2$? [MP PMT 1997]
 - (a) *Ca*
- (b) Sr
- (c) *Ba*
- (d) Ra

Point out the incorrect statement regarding Be 63. (Group-IIA)

[MP PMT 1997]

- (a) It forms an ionic carbide
- (b) Its carbonate decomposes on heating
- (c) Its halides are covalent
- (d) It is easily attacked by water
- Beryllium differs from rest of the members of its family (Group-IIA) in many ways. The reason for this is its

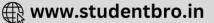
[MP PMT 1997]

- (a) Small size and higher electronegativity
- (b) Small size and lower electronegativity
- (c) Large size and lower ionisation energy
- (d) Large size and largest ionic radius
- The oxide, which is best soluble in H_2O is [BHU 2001]
 - (a) $Ba(OH)_2$
- (b) $Mg(OH)_2$
- (c) $Sr(OH)_2$
- (d) $Ca(OH)_2$
- The property of the alkaline earth metals that increases with their atomic number is [BHU 2001]
 - (a) Ionisation energy
 - (b) Electronegativity
 - (c) Solubility of their sulphates
 - (d) Solubility of their hydroxides
- In the Alkaline earth metals, the element forming predominantly covalent compound is [BHU 2001]
 - (a) Be
- (b) Mg

- (d) Ca
- 68. A mixture of lime paste is sand, water and [RPMT 1997]
 - (a) Gypsum
- (b) Slacked lime
- (c) Quick lime
- (d) Lime stone
- The formula for calcium chlorite is [CBSE PMT 1994, 96] (b) $Ca(ClO_3)$,
 - (a) $Ca(ClO_4)_2$
- (c) CaClO₂
- (d) $Ca(ClO_2)_2$
- Which pair of substances gives same gaseous product, when these 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







	766 s and p-Bloc	k Elements					
71.	Magnesium does not de	ecompose the	[AFMC 1999]		(c) $BeCl_2 < BaCl_2 < MgCl_2$	$Cl_2 < CaCl_2$	
	(a) Steam	(b) Hot wa	iter		(d) $BaCl_2 < CaCl_2 < MgC$	$Cl_2 < BeCl_2$	
	(c) Cold Water	(d) Semi h		83.	$MgCl_2.6H_2O$ when hea		[CPMT 1997]
72.	Alkaline earth metal		er than alkali	J	(a) Magnesium okychlo	_	2 22/2
	metals because metalli (a) Stronger	(b) Weake	r		(b) Magnesium dichlor		
	(c) Not present	(d) Volatil			(c) Magnesium oxide		
73.	Property of the all				(d) Magnesium chlorid	e	
, 3.	increases with their at			84.	Which of the followin	g hydroxide i	s insoluble in
	(a) Ionisation energy				water		
	(b) Solubility of their l	nydroxides					[AIIMS 2001]
	(c) Solubility of their s	sulphates			(a) $Be(OH)_2$	(b) $Mg(OH)$	2
	(d) Electronegativity				(c) <i>Ca(OH)</i> ₂	(d) <i>Ba(OH)</i> ₂	2
74.	A metal is burnt in air		on moistening	85-	Which of the following	statements is	false[BHU 2005]
	smells of NH_3 . The me			Ĺ	(a) $CaOCl_2$ gives OH^- ,		
	(a) <i>Na</i>	(b) Fe		solu	_		-
	(c) Mg	(d) <i>Al</i>			(b) Diamond and graph	ite are allotro	ops of carbon
75.	Alkaline earth metals ((c) Bleaching action o	f Cl ₂ in mois	t condition is
elem	(a) Halogens ients	(b) Repres	entative		not permanent		
	(c) Transition element	s (d) Inner	transition		(d) Calomel is Hg_2Cl_2		
elem	ients			86.	3		•
	(e) None of these				is water-soluble. It for		
76.	Which of the follow				becomes inert on hear hydroxide $M(OH)_2$ w		
	hydroxides is the stron		[CPMT 1996]		solution. Then M is	[AIEEE 2002	
	(a) $Be(OH)_2$	(b) <i>Mg(OH</i>	_		(a) <i>Mg</i>	(b) <i>Ba</i>	-
	(c) $Ca(OH)_2$	(d) <i>Ba(OH)</i>	O_2		(c) <i>Ca</i>	(d) <i>Be</i>	
77•	Which one of the follow	wing is the st	-	87.	In the lime (kiln), the r	• •	
	(a) n (ou)		[Pb. PMT 1998]	٥,٠	$CaCO_3(s) \rightarrow CO_2(g)$ goes		n because
	(a) $Be(OH)_2$	(b) $Mg(OH)$	<i>/</i> =		caeco 3(a) 7 co 2(g) good		
	(c) $Al(OH)_3$	(d) $Si(OH)$			(a) Of high temperatur		(Engg.) 2002]
78.	Lime stone is		[RPMT 1997]		(b) <i>CaO</i> is more stable		
	(a) CaO	(b) <i>Ca(OH)</i>	0_{2}			9	
	(c) Both (a) and (b)	(d) None o			(c) CO_2 escapes simul	taneously	
79.	Which of the alkaline	e earth meta	C		(d) CaO is not dissociate	ted	
	reducing agent	(b) <i>Sr</i>	[MP PMT 1995]	88.	The ionic compound <i>E</i>	<i>BaSO</i> ₄ is insol	uble in water
	(a) <i>Ca</i> (c) <i>Ba</i>	(d) Mg			due to		
00		_	[CDWT 1004]				[CPMT 1999]
80.	Plaster of paris harden (a) Giving off CO_2	-	[CPMT 1994]		(a) High lattice energy		-
			ng into CaCO ₃		(c) Low hydration ener		oth (a) and (c)
0.4	(c) Uniting with water	_		89.	which is used to reduce	-	
81.	Which is not soluble in		[CPMT 1994]		(a) Calcium hydroxide		=
	(a) $CaCO_3$	(b) $BaCO_3$			(c) Ammonium nitrate		ium chloride
	(c) $SrCO_3$	(d) All of t	hese	90.	Alkaline earth metals b	_	KCET (Med.) 2001]
82.	The correct order of th	e increasing	ionic character		(a) s – block in period:		
	is				(b) p – block in period	ic table	

[AFMC 2002]

(c) d – block in periodic table

(d) f – block in periodic table

CLICK HERE

91. The element having atomic number 56 belongs to

(a) $BeCl_2 < MgCl_2 < CaCl_2 < BaCl_2$

(b) $BeCl_2 < MgCl_2 < BaCl_2 < CaCl_2$

[MNR 1991; AFMC 1998]

is

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 MgCO₃, CaCO₃, BaCO₃ (a) In surgery and dentistry SrCO₃ 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 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 **106.** *Li* shows the diagonal relationship with [Pb.CET 2001] flame (b) B (a) Mq [AFMC 2001] (d) C (c) Al (a) Barium (b) Calcium **107.** A sodium salt on treatment with MgCl, 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) $NH_{\perp}Cl$ (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.** *Mq* burns in *CO* to produce [Pb.PMT 2001] (a) Mg (b) Sr (a) MgO_2 (b) $MgCO_3$ (c) Ca (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) $CaSiO_3.MgCO_3$ combines with oxygen [MH CET 2000]

[DCE 2003]

(d) $MgCl_2.5MgO.xH_2O$

(a) $Ca[B_3O_4(OH)_2].2H_2O$

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

(c) $Ca(OH)_2$

111. Colemnite is



[AFMC 2004]

(b) CaO

102. Slow acting nitrogenous fertilizer among the

(d) Ca_2O_2

(a) Ca

(c) *CaO*₂

following is

94.

95.

(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_3 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
- 6. Identify the statement that is not correct as far as 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 $30^{\circ}C$ 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
- 9. Which metal burn in air at high temperature with 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
- 14. 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 [Kerala PMT 2004]
 - (a) Be_2C
- (b) Graphite
- (c) Titanium
- (d) SiC
- (e) B_AC



Which of the following is known as inorganic 20. henzene

[Pb. CET 2001]

- (a) Borazine
- (b) Boron nitride
- (c) *p*-dichlorobenzene
- (d) Phosphonitrilic acid
- Which of the following is only acidic in nature 21.

[AIIMS 2004]

- (a) $Be(OH)_2$
- (b) $Mg(OH)_2$
- (c) $B(OH)_3$
- (d) $Al(OH)_3$
- 22. Moissan boron is

[DCE 2003]

- (a) Amorphous boron of ultra purity
 - (b) Crystalline boron of ultra purity
 - (c) Amorphous boron of low purity
 - (d) Crystalline boron of low purity
- Which of the following does not exist in free form 23.

[Kerala PMT 2004]

- (a) BF_3
- (b) BCl_3
- (c) BBr_3
- (d) BH_3
- (e) None of these
- 24. Alumina is

[DCE 2002]

- (a) Acidic (c) Amphoteric
- (b) Basic
- (d) None of these
- The most abundant metal in the earth crust is 25.

[Pb. CET 2004]

(a) Al

(b) Ca

(c) Fe

- (d) Na
- 26. Crystalline metal can be transformed into metallic glass by

[NCERT 1984]

- (a) Alloying
- (b) Pressing into thin plates
- (c) Slow cooling of molten metal
- (d) Very rapid cooling of a spray of the molten metal
- Which metal is protected by a layer of its own 27. oxide

[NCERT 1981; DPMT 1983; BHU 1998]

- (a) *Al*
- (b) Ag
- (c) Au
- (d) Fe
- 28. Aluminium is a self-preserving metal, because
 - (a) It is not tarnished by air
 - (b) A thin film of basic carbonate on its surface
- (c) A non-porous layer of oxide is formed on its surface
 - (d) It is not affected by salt water
- Anhydrous AlCl₃ cannot be obtained from which 29. of the following reactions [CPMT 1987]
 - (a) Heating $AlCl_3.6H_2O$

- (b) By passing dry HCl over hot aluminium powder
- (c) By passing dry Cl_2 over hot aluminium powder
 - (d) By passing dry Cl2 over a hot mixture of alumina and coke
- An element A dissolves both in acid and alkali. It 30. is an example of [NCERT 1972]
 - (a) Allotropic nature of A(b)Dimorphic nature of A
 - (c) Amorphous nature of A (d)Amphoteric nature of A
- Hydrogen gas will not reduce

[IIT 1984]

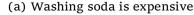
- (a) Heated cupric oxide
- (b) Heated ferric oxide
- (c) Heated stannic oxide
- (d) Heated aluminium oxide
- 32. Conc. HNO_3
 - (a) Reacts with aluminium vigrously
- (b) Reacts with aluminium to form aluminium nitrate
 - (c) Does not react with aluminium
 - (d) Reacts with platinum
- Anhydrous AlCl₃ is obtained from

[BHU 1980; CPMT 1982]

- (a) HCl and aluminium metal
- (b) Aluminium and chlorine gas
- (c) Hydrogen chloride gas and aluminium metal
- (d) None of the above
- Which is true for an element R present in III 34. group of the periodic table [EAMCET 1991]
 - (a) It is gas at room temperature
 - (b) It has oxidation state of +4
 - (c) It forms R_2O_3
 - (d) It forms RX_2
- When Al is added to KOH solution 35.

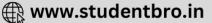
[NCERT 1974, 76; CPMT 1977]

- (a) No action takes place
- (b) Oxygen is evolved
- (c) Water is produced
- (d) Hydrogen is evolved
- Aluminium is more reactive than iron. But aluminium is less easily corroded than iron [KCET 1993] because
 - (a) Aluminium is a noble metal
 - (b) Oxygen forms a protective oxide layer
 - (c) Iron undergoes reaction easily with water
 - (d) Iron forms mono and divalent ions
- Aluminium vessels should not be washed with materials containing washing soda since[KCET 1993]









- (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 AlCl₃ 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.** AlCl₃ is [AFMC 1995]
 - (a) Anhydrous and covalent (b)Anhydrous and ionic
- (c) Covalent and basic (d) Coordinate and acidic
- 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]

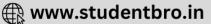
- (a) 3 parts of Al_2O_3 and 4 parts of Al
- (b) 3 parts of Fe_2O_3 and 2 parts of Al
- (c) 3 parts of Fe_2O_3 and 1 part of Al
- (d) 1 part of Fe_2O_3 and 1 part of Al
- **51.** Bauxite containing impurities of iron oxide is purified by

[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 NaOH solution at $50^{\circ}C$
 - (b) Bauxite ore is fused with Na_2CO_3
 - (c) Bauxite ore is fused with coke and heated at $1800^{\circ}C$ in a current of nitrogen
 - (d) Bauxite ore is heated with NaHCO₃
- **53.** Which one is used as a bye-product in Serpeck's process
 - (a) NH_3
- (b) *CO*₂
- (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
- In the electrolytic extraction of aluminium, 55. 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
- 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
- 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
- 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
- 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
- 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
- (c) For (i) Serpeck's process; for (ii) Baeyer's process
 - (d) For (i) Baeyer's process; for (ii) Serpeck's process
- 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
- In the commercial electrochemical process for 63. 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 $Al(OH)_3$
- In electrolysis of aluminium oxide which of the 64. following is added to accelerate the process[AFMC 1999; C
 - (a) Silica
- (b) Cryolite
- (c) Nickel
- (d) Silicate
- 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
- 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
- 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
- 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
- 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
- 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

	Bottom Layer	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 <i>Al</i>	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

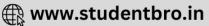
[CBSE PMT 2005]

- (a) B_2H_6
- (b) C_2H_6
- (c) PH₃
- (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 $Pb(OH)_2$
- (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

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 (σ) and one $pi(\pi)$ bond
 - (b) One sigma (σ) and two pi (π) bond
 - (c) One sigma (σ) and half pi (π) bond
 - (d) One sigma (σ) bond
- **13.** Metalloid among the following is **[DPMT 2001]**
 - (a) *Si*

(b) C

[Pb-PMA-2000]

(d) *Ge*

14. 'Lead pencil' contains

[DPMT 2001; IIT 1990]

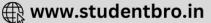
- (a) *PbS*
- (b) Graphite
- (c) FeS
- (d) *Pb*
- **15.** Nitrogen gas is absorbed by
 - d by [DPMT 2001]
 - (a) Calcium hydroxide
- (b) Ferrous sulphate
- (c) Calcium carbide
- (d) Aluminium carbide
- **16.** In laboratory silicon can be prepared by the reaction

[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



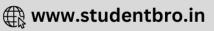




772 s and p-Block Elements (c) Covalent and ionic tendency Which gas is used in excess water 29. [BVP 2003] (d) Different valency (a) *CO*₂ (b) SO, Colour is imported to glass by mixing [Pb. PMT 2002] (c) CO (d) Water vapours The compound which does not possess a peroxide (a) Synthetic dyes (b) Metal oxide 30. linkage is (c) Oxides of non-metal (d) Coloured salt (b) CrO₅ (a) Na_2O_2 Which of the following is insoluble in water [MP PET 2002] (d) PbO_2 (a) Na_2CO_3 (b) CaCO₃ (c) H_2SO_5 Silicon is an important constituent of [MH CET 2001] 31. (c) $ZnCO_3$ (d) $Al_2(CO_3)_3$ (a) Rocks (b) Amalgams 20. In which of the following the inert pair effect is (c) Chlorophyll (d) Haemoglobin most prominent [MP PMT 2000] Carborundum is 32. (a) C (b) Si [AFMC 2002; MH CET 2003; BHU 2003, 05] (c) Ge (d) Pb (a) SiC (b) AlCl₃ 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 4 sol SiF_4 gets hydrolysed giving [Orissa JEE 2002] 33. Which of the following glass is used in making (a) SiO_2 (b) $Si(OH)_2F_2$ 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_{\perp} (b) H_2SiF_6 (c) H_2SiO_3 (d) Na_3AlF_6 (d) Polymeric mixture The type of glass used in making lenses and 24. H_2O_2 on reaction with *PbS* gives 35. [RPET 2003] prisms is (a) *PbO* (b) $PbSO_4$ [JIPMER 1999] (c) PbO_2 (d) PbHSO 4 (a) A flint glass (b) Jena glass (c) Pyrex glass (d) Quartz glass Soldiers of Napolean army while at Alps during 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 (b) NaHCO₃ (a) Na_2CO_3 powder. This transformation is related to (c) H-COONa(d) CH₃COONa [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 low to temperatures gives (d) An interaction with water vapour contained in [Roorkee 2000] the humid air (a) CO only (b) CO2 only 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. zeolites is false (b) Adding more lead sulphate into reverberatory 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

			s and p-Bloc	k Elements 773
	(c) They are used as cation exchangers	50.	Red lead is [C	PMT 1972, 74, 94; MNR 1985;
	(d) They have open structure which enables them			ihar CEE 1995; MP PET 1995]
	to take up small molecules	_	(a) Pb_3O_4	(b) <i>PbO</i>
39.	Which of the following cuts ultraviolet rays[AFMC 200	04]	(c) PbO_2	(d) Pb_4O_3
	(a) Soda glass (b) Crooke's glass	51.	White lead is [CPM	T 1983, 93, 2002; MNR 1984;
40	(c) Pyrax (d) None of these In IIIA group, <i>Tl</i> (thalium) shows +1 oxidation		MP PMT 19	95; UPSEAT 1999; DCE 2000]
40.	state while other members show +3 oxidation		(a) $PbCO_3$	(b) PbCO ₃ .PbO
	state. Why		(c) $2PbCO_3.Pb(OH)_2$	(d) $2PbSO_4.PbO$
	[JEE Orissa 2004]	52.	Lead pipes are corroded	l quickly by [AFMC 1981]
	(a) Presence of lone pair of electron in Tl		(a) Dil. H_2SO_4	(b) Conc. H_2SO_4
	(b) Inert pair effect		(c) Acetic acid	(d) Water
	(c) Large ionic radius of Tl ion	53.	In silicon dioxide	[AIEEE 2005]
	(d) None of these			is surrounded by four
41.	Carbon suboxide C_3O_2 has [DCE 2003]		oxygen atoms an	d each oxygen atom is
	(a) Linear structure		bonded to two silic	
	(b) Bent structure			is surrounded by two
	(c) Trigonal planar structure		oxygen atoms an bounded to two sili	d each oxygen atom is
	(d) Distorted tetrahedral structure	_		led to two oxygen atoms
42.	Which of the following is a mixed oxide[Pb. CET 2003]]		onds between silicon and
	(a) Fe_2O_3 (b) PbO_2		oxygen atoms	onds between sincon und
	(c) Pb_3O_4 (d) BaO_2	54.	Litharge is chemically	[DPMT 1984; JIPMER 2001]
43.	Noble gases are absorbed on [BVP 2004]		(a) <i>PbO</i>	(b) <i>PbO</i> ₂
	(a) Anhydrous <i>CaCl</i> ₂ (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.		nfiguration is of group
	(a) Ferrous sulphate (b) Copper sulphate		(a) IV	(b) III
	(c) Sodium alumino silicate (d) Zinc sulphate	-6	(C) V	(d) II compounds of elements in
45.	Which of the following statement is correct with	50.	· ·	xpect to be most ionic in
	respect to the property of elements in the carbon		character	
	family with an increase in atomic number, their [Pb. (a) Atomic size decreases	CET 20	02]	[NCERT 1978]
	(a) Atomic size decreases(b) Ionization energy increases		(a) CCl ₄	(b) SiCl ₄
	(c) Metallic character decreases		(c) PbCl ₂	(d) <i>PbCl</i> ₄
	(d) Stability of +2 oxidation state increases	57•	Which of the following	compounds of lead is used
46.	When tin is treated with concentrated nitric acid[DCI			•
_	(a) It is converted into stannous nitrate	-	(a) <i>PbO</i>	(b) <i>PbO</i> ₂
	(b) It is converted into stannic nitrate		(c) <i>PbCl</i> ₂	(d) None of these
	(c) It is converted into metastannic acid	58.	Type metal is an alloy of	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 th	e three metals
	(a) $Pb + Zn + Sn$ (b) $Pb + Zn$		(b) More amount of lead	
40	(c) $Pb + Sn$ (d) $Sn + Zn$		(c) More amount of ant	imony
48.	A metal used in storage batteries is (a) Copper (b) Lead		(d) More amount of tin	
	(c) Tin (d) Nickel	59.	Which is correct oxidati	ion state of lead[AFMC 1987]
49.	Name of the structure of silicates in which three		(a) + 2, + 4	(b) + 1, + 2
	oxygen atoms of $[SiO_4]^{4-}$ are shared is [IIT 2005]		(c) $+ 3, + 4$	(d) + 4
	(a) Pyrosilicate	60.	Sugar of lead is	
	(b) Sheet silicate		(a) 2 <i>PbSO</i> ₄ . <i>PbO</i>	(b) $(CH_3COO)_2Pb$
	(c) Linear chain silicate		(c) $PbCO_3$	(d) $PbCO_3.Pb(OH)_2$
	(d) Three dimensional silicate			





61.	Which of the follow	ving compounds has peroxide		Pb. PMT 2000; AFMC 19	88, 92; MP PE	Γ 1997, 2000, 01]
	linkage			(a) N_2O	(b) <i>NO</i>	
	(a) N O	[CPMT 1988]		(c) N_2O_3	(d) N_2O_5	
	(a) Pb_2O_3	(b) SiO ₂	4.	Metaphosphoric acid	has the formu	ıla[CPMT 1973, 89, 93]
	(c) <i>CO</i> ₂	(d) PbO_2	•	(a) H_3PO_4	(b) <i>HPO</i> ₃	1 0,0, 0,001
62.	Percentage of lead in	-		(c) H_2PO_3	(d) H_3PO	
	(a) 7ono	[CBSE PMT 1999]				-
	(a) Zero (c) 80	(b) 20 (d) 70	5.	Which of the following agent for ammonia ga	-	suitable drying
63.	Which of the follow			=		989; DPMT 1982;
٠,٠	Willest of the follows	[CPMT 1996]				J 1986, 96; 2001;
	(a) Fe	(b) <i>Cu</i>		0111111		Screening) 2000]
	(c) B	(d) <i>Pb</i>		(a) Calcium oxide		g , 1
64.	Red lead in an exam	ple of a/an oxide		(b) Anhydrous calciu	m chloride	
		[JIPMER 2001]		(c) Phosporus pentox		
	(a) Basic	(b) Super		(d) Conc. sulphuric a		
6-	(c) Mixed	(d) Amphoteric	6.	Each of the following		white and red
65.	which of the follows	ing lead oxides is 'Sindhur' [MP PET 2002]	٠.	phosphorus except th	at they	wifice difd red
	(a) <i>PbO</i>	(b) <i>PbO</i> ₂		(a) Are both soluble i	in CS_2	
	(c) Pb_2O_3	(d) Pb_3O_4		(b) Can be oxidised b	y heating in a	ir
66.	Element showing th	e phenomenon of allotropy is		(c) Consists of same	kind of atoms	
	O	[MP PMT 1999]		(d) Can be converted	into one anot	her
	(a) Aluminium	(b) Tin	7•	Which of the following	_	
	(c) Lead	(d) Copper		(a) Orthophosphorus		= =
67.	Which of the follow	ing element is a metalloid		(c) Metaphosphoric a		=
		[CPMT 2004]	8.	Phosphine is prepare	d by the react	ion of
	(a) Bi	(b) <i>Sn</i>				P PET/PMT 1988]
	(c) Ge	(d) <i>C</i>		(a) P and H_2SO_4	(b) <i>P</i> and	d NaOH
68.	Which gas is liberat	ed when Al_4C_3 is hydrolysed		(c) P and H_2S	(d) <i>P</i> and	$d HNO_3$
		[AFMC 2005]	9.	Which of the following	ng is not know	m[MP PET/PMT 1988 ;
	(a) CH_4	(b) C_2H_2			CBSE PMT 198	9; MP PET 1993]
	(c) C_2H_6	(d) CO_2		(a) NCl_5	(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	the phosphor	rus molecule is
	(a) HCl	(b) HF		[CPMT	1976, 80, 84, 9	o; BHU 1984, 86;
	(c) HI	(d) HBr				NCERT 1977]
	NI'd			(a) <i>P</i>	(b) P_4	
	Nitro	gen family		(c) P_2	(d) P_5	
1.		wing elements does not form	11.	White phosphorus (P	• ,	[IIT 1998]
	stable diatomic mole (a) Iodine	ecules (b) Phosphorus	[CBSP FMT^P1589 ; in ple bo (b) Four <i>P - P</i> single l		
	(c) Nitrogen	(d) Oxygen		(c) Four lone pairs of	f electrons	
2.	=	xture of[DPMT 1982; CPMT 1978]		(d) <i>PPP</i> angle of 60°		
۳.	(a) CO and N_2	(b) CO_2 and H_2	12.	Ammonium nitrate de	-	•
	(c) CO and H_2	(d) CO_2 and N_2		[NCERT :	1974,75; CPMT	1973, 78, 88, 94; AMU 1984]
3.	Which one of the fo	ollowing combines with $Fe(II)$		(a) Ammonia and nit	ric acid	



(b) Nitrous oxide and water

(c) Nitrogen, hydrogen and ozone

ions to form a brown complex

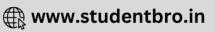
[AIIMS 1982, 83, 87; BHU 1998; CBSE PMT 2000;

					Siden Elements //5
		en dioxide and hydrogen	23.	On heating a mixtur	re of NH_4Cl and KNO_2 we get
13.	In Birkeland-Eyde procis	ess, the raw material used			[CPMT 1972, 79; NCERT 1977]
	15	[CPMT 1982, 86]		(a) NH_4NO_3	(b) N_2
	(a) Air	(b) NH_3		(c) N_2O	(d) <i>NO</i>
	(c) NO ₂	(d) HNO ₃	24.	Which of the folloanhydride of <i>HNO</i> ₃	wing oxide of nitrogen is the
l 4.	nitrate and Ammoniu	itrates, Lead nitrate, Silver m nitrate; the one that aving any solid residue is[NC	ERT 198	CRSE DMT 1080 0	7; MP PET/PMT 1988; KCET 1991; 1,99; EAMCET 1991; NCERT 1975; MP PET 1989; MP PMT 1994]
	(a) Lead nitrate	(b) Ammonium nitrate		(a) <i>NO</i>	(b) N_2O_3
	(c) Silver nitrate	(d) Sodium nitrate			
5.	which is most reactive is [CPMT 1983; NCERT 1978; CBSE PMT 1999; Kurukshetra CEE 1998]		25.	_	(d) N_2O_5 unufactured by heating in a nixture of NCERT 1977; CPMT 1974,
	(a) Violet phosphorus	(b) Scarlet phosphorus		(a) Bone ash and co	oke
	(c) Red phosphorus	(d) White phosphorus		(b) Bone ash and si	lica
6.	Phosphine is generally	prepared in the laboratory		(c) Bone ash, silica	and coke
		[CPMT 1983, 2003]		(d) None of these	
ydr	ogen	phorus in a current of	26.	dissolved in water	orms a solid oxide which when forms an acidic solution, the
	solution of caustic	phosphorus with aqueous		element is	[CPMT 1972, 78]
	(c) By decomposition o			(a) Argon	(b) Potassium
				(c) Phosphorus	(d) Sulphur
	solution of caustic s	osphorus with an aqueous soda	27.	-	PO_4 occurs in following stages
7.		elements is most metallic		3	[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	v many oxides
3.	The basicity of orthoph	osphoric acid is[CPMT 1984, 9)1]	(a) 3	(b) 4
	(a) 2	(p) 3		(c) 5	(d) 6
	(c) 4	(d) 5	29.	The P-P-P bond ang	le in white phosphorus is
).	HNO ₂ acts as	[AFMC 1992]			[MP PET 1991]
	(a) Oxidising agent	(b) Reducing agent		(a) 120°	(b) 109 ° 28′
	(c) Both (a) and (b)	(d) Its solution is stable		(c) 90°	(d) 60°
0.	[CPM	t be obtained by heating T 1989; IIT 1985; CPMT 1993]	30.		nate on heating gives 8; CBSE PMT 1993; MP PMT 1993] e and ammonia
	(a) KNO_3	(b) $Pb(NO_3)_2$		(b) Chromic acid an	
	(c) $Cu(NO_3)_2$	(d) $AgNO_3$		(c) Chromium oxide	e and nitrogen
l.	When heated NH_3 i evolved is	s passed over <i>CuO</i> gas	31.		d nitric acid is heated, it
		[BCECE 2005]		decomposes to give (a) O_2 and N_2	(b) <i>NO</i>
	(a) N_2	(b) N_2O		(c) N_2O_5	(d) NO_2 and O_2
	(c) HNO_3	(d) NO_2			
2.	Non-combustible hydric	de is [CPMT 1979]	32.		catches fire in air at 30° C and er is [BHU 1973; MP PET 1989,99]
	(a) NH_3	(b) <i>PH</i> ₃		(a) Calcium	(b) Sodium
	(c) AsH_3	(d) SbH 3		(c) Phosphorus	(d) Zinc
	(-) 110113	(, 5011 3	33.	A 7 11 C	nia in water contains

	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,							
	(a) H ⁺	[CBSE PMT 1991]	gyps	(b) A mixtu	ire of prim	ary calcium	phospha	ate and
	(b) <i>OH</i> -		871-	(c) Normal	calcium ph	osphate		
				(d) Soluble	_	-		
	(c) Only NH ⁺ ₄		44.	If phospho	-	-	to reac	t with
	(d) OH^- , NH_4^+ and NH			sufficient q	uantity of <i>I</i>	<i>NaOH</i> , the p	roduct ol	btained
34.		V A group (N, P, As, Sb and		is		IDDMT 400	o. MD DM	т 1000]
		wing properties shows an from nitrogen to bismuth[C	DMT 109	8 ኅ ኬ) <i>NaHDO</i>		(b) Na_2H		1 1983]
	(a) Stability of +3 oxida		rwii ige	5		=	· ·	
	(b) Reducing character			(c) <i>NaH</i> ₂ <i>PO</i>	·	(d) Na_3Pa_3	O_4	
	(c) Electronegativity	•	45.	White phosp	_			_
	(d) Acidic nature of the	pentoxide				T (Med.) 200		T 1990]
35∙	The important method:	for the fixation of nitrogen		(a) P_5 mole		(b) P_4 m		
	is			(c) P_6 mole	ecules	(d) P_2 m	olecules	
	(a) Hahan	[CPMT 1985, 94]	46.	In the catal	•			
	(a) Haber(c) Deacon	(b) Solvay(d) Fischer method		formed whi		n the prepa	ration of	HNO_3 .
36.	Which of the following			This oxide i				
50.	winds of the following	[AFMC 1987; KCET 1991]		(-) N 0	[CPMT 1	1984; KCET 1	990; AIIM	S 1996]
	(a) White phosphorus	(b) <i>CH</i> ₄		(a) N_2O_5		(b) N_2O_4		
	(c) H_2O	(d) NaCl		(c) NO_2		(d) <i>NO</i>		
37.	-	en is prepared by heating	47.	Nitric acid o	oxidises pho	_		
3/•	(a) Calcium cyanamide					[CPMT 198		R 2002]
	•	de (d)Ammonium nitrite		(a) $H_2P_2O_7$		(b) H_3PC	-	
38.	Nitrous oxide			(c) P_2O_5		(d) H_3PC	4	
_	(a) Is a mixed oxide		48.	Which one	of the follo	wing staten	nents is t	rue for
	(b) Is an acidic oxide			HNO_2				
	(c) Is highly soluble in	hot water					[CPMT 19	80, 84]
	(d) Supports the combu	stion of sulphur		(a) It is ver	-	_		
39.	Which of the following	represents laughing gas		(b) It cann	ot act botl	n as an ox	idant an	d as a
		pal MEE 1995;MP PMT 1990;	reau	ctant	ot act ac an	ovidicina oc	ont	
		95; RPMT 1999; AFMC 2002]		(c) It canno (d) It canno				
	(a) NO	(b) N_2O	49.	Which oxide			[MP PE	T 1000]
	(c) NO_2	(d) N_2O_3	49.	(a) P_2O_3		(b) Bi_2O_3	_	1 1990]
40.	NO_2 is a mixed oxide	is proved by the first that						
	with NaOH, it forms			(c) As_2O_3		(d) B_2O_3		
	(a) Nitrites salt		50.	Which acid	is formed b	- 2 3		T 1991]
	(b) Nitrates salt			(a) H_3PO_4		(b) H_3PC	3	
	(c) Mixture of nitrate a	nd nitrite		(c) HPO_3		(d) $H_4 P_2$	O_7	
	(d) Ammonia		51.	Which nitro	gen trihalio	des is least l	oasic	
41.	oxide with dil. HNO ₃	g metal produces nitrous		TII]	1987; Kuruk	shetra CEE 1	998; CPM	T 1999]
		(h) 7		(a) NF_3		(b) NCl_3		
	(a) Fe	(b) Zn		(c) NBr_3		(d) NI_3		
	(c) Cu	(d) Ag	52.	Dehydrated	phosphoru	s trichloride	e in water	r gives
42.	form	g acid exist in polymeric	_	•	• •			T 1990]
	(a) HPO_3	(b) $H_4 P_2 O_7$		(a) HPO_3		(b) H_3PC	4	
	-			(c) H_3PO_2		(d) H_3PC)3	
	(c) H_3PO_4	(d) None of these	53.	Which is	used in			of safe
43.	Superphosphate of lime		55.	matchsticks				
gyng		al calcium phosphate and				[DPMT 1982		974, 75]
gyps	· UIII			(a) White p	=	_		
				(c) Red pho	sphorus	(d) Selen	ıum	

54.	Which oxide of nitroger			(d) <i>CaC</i> ₂	
		1987; Kurukshetra CEE 1998]	66.	On adding water to	$BiCl_3$ solution in HCl , the
	(a) N_2O	(b) <i>NO</i>		compound produced	is
	(c) N_2O_5	(d) NO_2		(a) Bi_2O_3	(b) $Bi(OH)_3$
55.	Which oxide do not act			(c) BiOCl	(d) BiOCl ₂
	(a) NO	[MP PET 1990] (b) NO_2	67.	V-A group precipitate	e was dissolved in HNO_3 and
	(c) N_2O	(d) N_2O_5			of NH_4OH . It gives a white
-6	-	ation number of nitrogen		ppt. because of	
56.	will be	ation number of introgen		(a) $Cu(OH)_2$	(b) $Cd(OH)_2$
	WIII DE	[MP PET 1990]		(c) $Bi(OH)_3$	(d) $Hg(OH)_2$
	(a) + 3	(b) + 5	68.	N, P, As, Sb, Bi eleme	ents belong to [DPMT 1982]
	(c) -3 and $+3$	(d) + 3 and + 5		(a) VA group	(b) IVA group
57•		the oxidation state of		(c) VIIA group	(d) VB group
	phosphorus is+4	[MP PET 1991]	69.		lowing elements occur free in
	(a) P_4O_{11}	(b) P_4O_8		nature	[CDWT 4000]
	(c) P_4O_6	(d) $H_3 PO_4$		(a) Nitrogen	[CPMT 1988] (b) Phosphorus
58.	. 0	the oxidation state of		(c) Arsenic	(d) Antimony
50.	nitrogen is -1		70.		ng 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) <i>P</i>	(d) As
59.	Which of the following		71.	Which does not form	_
		P PMT 1990; CBSE PMT 1996]		(a) <i>N</i> (c) <i>As</i>	(b) <i>P</i> (d) <i>Bi</i>
	(a) $P_4 O_6$	(b) $P_4 O_{10}$	72.	The strongest base is	
_	(c) As_4O_6	(d) As_4O_{10}	/=-	_	9; CPMT 1997; MP PET 2001,03]
60.		f hydrides of the V-group he order [CBSE PMT 1996]		(a) NH_3	(b) PH_3
	(a) $SbH_3 > PH_3 > AsH_3$			(c) AsH_3	(d) SbH_3
	(b) $NH_3 > SbH_3 > PH_3 >$		73.	The most stable hydr	ride is [EAMCET 1988]
	(c) $NH_3 > PH_3 > AsH_3 >$			(a) NH_3	(b) PH_3
	(d) $SbH_3 > AsH_3 > PH_3$	-		(c) AsH_3	(d) SbH_3
61.	Which is least stable	-	74.	Which has the lowest	t boiling point[CBSE PMT 1989]
01.	(a) BiH_3	(b) SbH ₃		(a) NH_3	(b) PH_3
	(c) AsH_3	(d) <i>PH</i> ₃		(c) AsH_3	(d) SbH_3
62.	9	is not hydrolysed[DPMT 2005	, 75·	Which is the most ex	plosive[BHU 1984; Roorkee 1989
02.	(a) $AsCl_3$	(b) PF_3)]		IIMS 1996; MP PMT 1985, 2001]
	(c) $SbCl_3$	(d) NF_3		(a) NCl_3	(b) PCl_3
C -		*	1	(c) $AsCl_3$	(d) All of these
63.		re is maximum for[MP PET 19	⁹⁰ / ₇ 6.		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 2005	77.	Of the following, non	n-existent compound is
	(a) White phosphorous(c) Potassium	(d) Calcium		(a) PH_4I	[NCERT 1975, 79]
65.		g substances is used as a		•	(b) As_2O_3
٠,٠	fertilizer	o cassimises is abea as a	~	(c) SbCl ₂	(d) As_2H_3
	(a) $K_2SO_4.Al_2(SO_4)_3.24$	H_2O	78.	Pure N_2 gas is obtain	
	(b) $Ca(H_3PO_4)_2H_2O + Co$	aSO ₄			(b) $NH_4Cl + NaNO_2$
	(c) NaAlO ₂			$(c) N_2O + Cu$	· = - ·
			79.	Pure nitrogen can be	prepared from





778 s and p-Block Elements [KCET 1991; AFMC 1993; AMU 1985] (a) CO (b) NO (a) $NH_{\perp}OH$ (b) Ca_3N_2 (c) ClO₂ (d) N_2O_5 (c) NH_4NO_2 (d) $Ba(NO_3)_2$ On strongly heating $Pb(NO_3)_2$ crystals, the gas 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) NaNO₂ (d) NaNO₃ (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 [AFMC 1990] (a) Fe (b) Cu The cyanide ion, CN^- and N_2 are isoelectronic. (d) Sn (c) Zn But in contrast to CN^- , N_2 is chemically inert 93. 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 (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. neutral (c) It is a typical non-metal (d) d-orbitals are available for bonding (a) N_2O_5 (b) N_2O_3 The element which is essential in nitrogen 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 (c) NO_2 (d) N_2O_5 **85.** Laughing gas is prepared by heating[EAMCET 1980] The reddish brown coloured gas formed when (a) $NH_{\perp}Cl$ (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 (a) NO and NO_2 (b) NO_2 and O_2 **87.** Which of the following is not correct for N_2O [CPMT 1984]

(a) It is called laughing gas

(b) It is nitrous oxide

(c) It is not a linear molecule

(d) It is least reactive in all oxides of nitrogen

88. Which of the following oxides of nitrogen is the anhydride of nitrous acid

(a) NO

(b) N_2O_3

(c) N_2O_4

high

82.

(d) N_2O_5

Which of the following is a true acidic anhydride

[NCERT 1977]

When lightning flash is produced, which gas may

Of the following, which has three electron bond in

[CPMT 1988]

(c) NO_2 and N_2O

(d) NO and O_2

Which of the nitrates on strong heating leaves the 99. [KCET 1990] metal as the residue

(a) $AgNO_3$

(b) $Pb(NO_3)_2$

(d) $Al(NO_3)_3$

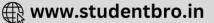
(c) $Cu(NO_3)_2$ [NCERT 1975; AIIMS 1991] **100.** Nitrogen dioxide

[KCET 1989]

(a) Dissolves in water forming nitric acid

(b) Does not dissolve in water





[KCET 1989] (c) Dissolves in water to form nitrous acid and gives off oxygen (a) Heating it with a catalyst in an inert (d) Dissolves in water to form a mixture of 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 (a) CO_2 and H_2O (d) Melting it and pouring the liquid into water (b) CO and H_2O 112. Bones glow in the dark because [EAMCET 1980] (c) CO, CO_2 and H_2O (d) Oxalic and acid (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 (a) N_2O (b) NO (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*₂ (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 (b) Phosphoric acid (a) Phosphine **108.** PCl_5 exists but NCl_5 does not because (c) Phosphorus acid (d) No reaction [EAMCET 1977, 82] 117. 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 (b) NH_3 (a) PH_3 (c) Black (d) All stable

(c) P_4O_6



(d) $P_4 O_{10}$

120. Phosphine is produced by adding water to [KCET 1991]

phosphorus by

111. Red phosphorus can be obtained from white

780 s and p-Block Elements (a) CaC_2 (b) HPO_3 (a) N_2 (b) HNO_3 (c) Ca_3P_2 (d) $P_4 O_{10}$ (c) NH_3 (d) PH_3 121. When aluminium phosphide is treated with dil. 132. The number of hydroxyl group in pyrophosphoric sulphuric acid acid is [KCET 1989] [KCET 1993] (a) SO_2 is liberated (b) PH_3 is evolved (a) 3(b) 4 (d) H_2 is evolved (c) 5 (d) 7 (c) H_2S is evolved 133. There is very little difference in acid strength in 122. With reference to protonic acids, which of the the series H_1PO_3 , H_3PO_3 , and H_3PO_2 because [KCET 199 following statements is correct (a) Phosphorus in these acids exists in different (a) PH_3 is more basic than NH_3 oxidation states (b) PH_3 is less basic than NH_3 (b) The hydrogen in these acids are not all (c) PH_3 is equally basic as NH_3 bounded to the phosphorus (d) PH_3 is amphoteric while NH_3 is basic (c) Phosphorus is not a highly electronegative element **123.** One of the acid listed below is formed from P_2O_3 (d) Phosphorus oxides are less basic and the rest are formed from P_2O_5 . The acid **134.** $BiCl_3$ on hydrolysis forms a white precipitate of [NCERT 1975] formed from phosphorus (III) oxide is [CPMT 1988] (a) HPO_3 (b) $H_4 P_2 O_7$ (a) Bismuthio acid (c) H_3PO_4 (d) H_3PO_3 (b) Bismuth oxychloride **124.** P_2O_5 is heated with water to give (c) Bismuth pentachloride [CBSE PMT 1991; DPMT 2000] (d) Bismuth hydroxide (a) Hypophosphorus acid (b)Orthophosphorus acid 135. At high temperature nitrogen combines with (c) Hypophosphoric acid (d) Orthophosphoric acid calcium carbide (CaC_2) to give 125. Hypophosphorus acid is[NCERT 1977; MP PMT 1992] [DPMT 1981, 85; AFMC 1998; MP PET 2000] (a) A tribasic acid (b) A dibasic acid (a) Calcium cyanide (b) Calcium cyanamide (c) A monobasic acid (d) Not acidic at all (c) Calcium carbonate (d) Calcium nitride **126.** PCl_3 reacts with water to form 136. Calcium cyanamide on treatment with steam under pressure gives ammonia and [EAMCET 1984, 88; CPM [KCET 1990; CBSE PMT 1991; CPMT 2003] (a) Calcium carbonate (b) Calcium hydroxide (a) PH_3 (b) H_3PO_3 , HCl(c) Calcium oxide (d) Calcium bicarbonate (c) POCl₃ (d) H_3PO_4 137. Which one has the highest percentage of nitrogen **127.** H_3PO_3 is [KCET 1991; CBSE PMT 1993; [CPMT 1977, 79, 94; NCERT 1981; MP PMT 1980] AIIMS 1996, MP PET 2001; RPET 2003] (a) A tribasic acid (b) A dibasic acid (a) Urea (b) Ammonium sulphate (c) Neutral (d) A monobasic acid (c) Ammonium nitrate (d) Calcium nitrate **128.** Oxidation state of + 1 for phosphorus is found in 138. Superphosphate of lime contains [CPMT 1984] [MP PMT 1991; MP PET 2001] (a) $Ca_3(PO_4)_2$ (b) CaHPO₄ (a) H_3PO_3 (b) H_3PO_4 (c) $Ca_3(PO_4)_2 + H_3PO_4$ (d) $Ca(H_2PO_4)_2$ (c) H_3PO_2 (d) $H_4 P_2 O_7$ 129. Which is not an acid salt [MNR 1989] 139. Thomas slag is [CPMT 1988] (a) NaH_2PO_3 (b) $NaH_{2}PO_{2}$ (a) $Ca_3(PO_4)_2$ (b) MnSiO₃ (c) $Na_3HP_2O_6$ (d) $Na_4P_2O_7$ (c) CaSiO₃ (d) FeSiO 3 **130.** By the action of hot conc. H_2SO_4 , phosphorus 140. When equal weights of the two fertilizers, urea changes to and ammonium sulphate are taken, urea contains[KCET 19 [Roorkee 1992] (a) Less nitrogen than ammonium sulphate (a) Phosphorus acid (b) As much nitrogen as ammonium sulphate (b) Orthophosphoric acid (c) Twice the amount of nitrogen present in (c) Metaphosphoric acid ammonium sulphate (d) Pyrophosphoric acid (d) More than twice the amount of nitrogen 131. Cyanamide process is used in the formation of present in ammonium sulphate [BHU 1995]

- **141.** Which statement is wrong for *NO*
- [DPMT 2005]
- (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)
$$O = N = 0$$
 (b) $O = 0$ (c) $O = 0$ (d) $O = 0$ (e) $O = 0$ (e) $O = 0$ (f) $O = 0$ (

(c)
$$O = N = O$$

$$O = O$$

$$O = O$$

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

- **143.** Blasting of TNT is done by mixing
 - (a) NH₄Cl
- (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]

- (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) PH₃
- (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*₄⁺
- (c) *PCl*₆
- (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]

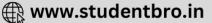
- (a) At the corners of tetrahedron
- (b) At the corners of a cube
- (c) At the corners of a four membered ring
- (d) At the centre and corners of an equilateral triangle
- 154. The most common minerals of phosphorus are

[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 PCl₅ 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
- $\textbf{159.} \ \ \, \text{Liquid ammonia is used for refrigeration because}$

[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_3) 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 HNO3
- (b) Zinc with concentrated HNO₃
- (c) Copper with cold and dilute HNO3
- (d) Heating NH_4NO_3
- **162.** What would happen when a solution of potassium chromate is treated with an excess of dilute nitric acid [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 PH_3

[BHU 2000]

- (a) PH_3 is not much stable
- (b) PH_3 is neutral towards litmus
- (c) PH_3 has fishy smell
- (d) PH3 is insoluble in water
- **166.** Nitrogen is obtained when NaNO, reacts with

[UPSEAT 2003]

- (a) NH_4Cl
- (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

[CPMT 2003]

- (a) $Na_2P_2O_7$
- (b) $Na_4P_2O_7$
- (c) $NaPO_4$
- (d) Na_2PO_2
- **170.** Which of the following is solid in nature

[UPSEAT 2003; AFMC 2004]

- (a) NO
- (b) $N_{2}O$
- (c) N_2O_3
- (d) N_2O_5
- 171. Which of the following is a cyclic phosphate

[KCET 1996]

- (a) $H_5 P_3 O_{10}$
- (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*
- [KCET 2002] (c) NCl₃
- (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) PCl_3 and HCl are formed and the mixture warms up
 - (c) *PCl*₅ 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.	When ammonia reacts product containing nitro	with sodium hypochlorite, ogen is [AFMC 2000]	189.	Which of the following soluble in ammonia	compounds is sparingly [JIPMER 1999]
	(a) N_2	(b) N_2O		(a) AgI	(b) AgBr
	(c) NH_2OH	2		(c) AgCl	(d) CuCl ₂
178	P_2O_5 is used extensively		190.		pes not leave a residue on
1/0.			190.	heating is	ies not leave a residae on
	(a) Reducing agent(c) Dehydrating agent			Ü	[JIPMER 1999; DCE 1999]
170		due to [DCE 2000; MP PET 200:	1]	(a) Na_2CO_3	(b) Ag_2CO_3
1/9•	(a) No vacant d -orbital	ade to [Bel 2000, MI 121 200.	- ,	(c) CuCO ₃	(d) $(NH_4)_2 CO_3$
	(b) High dissociation en	nergy	191.		g has the highest dipole
	(c) High electronegative	= -		moment	
	(d) None	9			[CBSE PMT 1997]
180.	Which show maximum	valency [CPMT 2003]		(a) NH_3	(b) PH_3
	(a) Phosphorus	(b) Tin		(c) SbH_3	(d) AsH_3
	_	(d) Bismuth	192.	The structural formula o	of hypophosphorus acid is
181.		Haber process for the			BCECE 2005; Pb. CET 2002]
	manufacture of NH_3	[MH CET 2001]		0	0
	(a) Pt	(b) $Fe + Mo$		(a) P_{QH}	(b) P OH
	(c) CuO	(d) Al_2O_3		H	OH
182.	_	mmonium hydroxide to a	_	(a) $H \stackrel{P}{\stackrel{ }{\nearrow}} OH$ MH CET 1999] (c) $H \stackrel{P}{{\nearrow}} OH$	0
	copper chloride solution		LI	MH CET 1999]	(d) (B)
	(a) A deep blue solution			$H \cap OH$	$OH \longrightarrow OOH$
	(b) No change is observ	of copper hydroxide is		11	OII
	obtained	or copper hydroxide is	193.	-	of the following fertilizers
	(d) Black precipitate of o	copper oxide is obtained			ty of the soil[CBSE PMT 1998]
183.	The product obtained	by heating $(NH_4)_2SO_4$ and		(a) Urea(c) Ammonium sulphate	(b) Potassium nitrate (d) Superphosphate of
	KCNO is		lime	_	(u) Super phosphate of
		[DPMT 2000]		Aqua-regia is	[Orissa JEE 2005]
	(a) Hydrocyanic acid			(a) 1:3 conc. HNO_3 and	- · · · · · · · · · · · · · · · · · · ·
.0.	(c) Ammonium cyanide			(b) 1: 2 conc. HNO_3 and	
104.		-O-P bonds in cyclic [IIT-JEE (Screening) 2000]		(c) $3:1$ conc. HNO_3 and	
	(a) Zero	(b) Two		(d) 2:1 conc. HNO_3 and	
	(c) Three	(d) Four	105	· ·	n is obtained on heating
185.		into the palm and washed	195.	ammonium nitrate at 25	_
	with water, it turns i	nto yellow. It shows the		(a) Nitric oxide	(b) Nitrous oxide
	presence of			(c) Nitrogen dioxide	(d) Dinitrogen oxide
		[CPMT 1997]	196.	The oxidation number of	• •
	(a) NO_2	(b) N_2O	5 - 0		shetra CEE 1998; DCE 2001]
	(c) <i>NO</i>	(d) N_2O_5		(a) -1 to +3	(b) -3 to +3
186.	Which of the following is	is nitrogenous fertilizers		(c) -3 to +5	(d) -5 to +1
		[CPMT 1999]	197.	Inorganic graphite is	[KCET 2003]
	(a) Bone meal	(b) Thomas meal		(a) $B_3 N_3 H_6$	(b) B_3N_3
40-	(c) Nitro phosphate	(d) Ammonium sulphate		(c) SiC	(d) $Fe(CO)_5$
187.	Which compound is rela	=	198.	Conc. HNO_3 can be store	d in container of[MH CET 2001]
	(a) <i>CO</i> ₂	[RPET 1999] (b) H ₂		(a) Al	(b) Sn
	(c) NO_2	(d) NH_3		(c) Cu	(d) <i>Zn</i>
100	-	•	199.	Which of the following c	_
100.	(a) Quick lime	[CPMT 2002; JIPMER 2002] (b) Slaked lime		(a) SbCl ₃	[JIPMER 1997] (b) <i>BiCl</i> ₅
	(c) Anhy. CaCl ₂	(d) None of these			
	(c) Anny. cuc ₂	(a) None of these		(c) SbCl ₅	(d) $AsCl_5$





	784 s and p-Block	Elements			
200.	The formula of an or H_3PO_4 . It is a	xyacid of phosphorus is	212.	One mole of calcium phosphide on reaction with excess water gives [IIT-JEE 1999]	
	[MP PMT 1996; (a) Dibasic acid (c) Tribasic acid	CPMT 1999; J & K CET 2005] (b) Monobasic acid (d) Tetrabasic acid		(a) One mole of phosphine(b) Two moles of phosphoric acid(c) Two moles of phosphine	
201.	(a) Na_2SO_4	fied as an acid salt[CPMT 199 (b) BiOCl		(d) One mole of phosphorous pentoxide On heating ammonium dichromate, the gas evolved is	
202.	(c) $Pb(OH)Cl$ In NH_3 and PH_3 , the co	(d) Na_2HPO_4 mmon is [AFMC 1995]		[IIT-JEE 1999] (a) Oxygen (b) Ammonia	
	(a) Odour(c) Basic natureWhich one of the fol stable	(b) Combustibility (d) None of these lowing hydrides is least	214.	(c) Nitrous oxide (d) Nitrogen In compounds of type ECl_3 , where $E = B$, P , As or Bi , the angles $Cl - E - Cl$ for different E are in the order	
	(a) AsH_3	(b) SbH ₃		[IIT-JEE 1999]	
204.	(c) NH ₃ Which element from V compound with hydroge	(d) PH ₃ group, gives most basic	215.	(a) $B > P = As = Bi$ (b) $B > P > As > Bi$ (c) $B < P = As = Bi$ (d) $B < P < As < Bi$ Appropriation reaction with hypochlorite anion,	
	(a) Nitrogen (c) Arsenic	(b) Bismuth (d) Phosphorus	ų 1	can form [IIT-JEE 1999]	
205.		vo series of salts is [KCET 199 (b) H_3PO_3	6]	(a) NO (b) NH_4Cl (c) N_2H_4 (d) HNO_2	
206.	(c) H_3BO_3 Which gas is obtained	(d) H_3PO_2 when urea is heated with	216.	Orthophosphoric acid represents the molaysis condition due to (a) Hydrogen bonding	
	HNO_2 (a) N_2	[CPMT 1996] (b) \boldsymbol{H}_2		(b) Phosphorous group(c) Maximum oxygen group(d) Tribasicity	
207.	(c) O_2 Atomic number of N is	(d) NH_3 7. The atomic number of	217.	Which of the following elements forms a strongly acidic oxide	
	IIIrd member of nitroger (a) 23 (c) 33	•	218.	(a) <i>P</i> (b) <i>As</i> (c) <i>Sb</i> (d) <i>Bi</i> In group V-A of the periodic table nitrogen forms only a trihalide but other elements form pentahalides also. The reason is	
	bond (a) PH_3	[CPMT 1996] (b) P_2H_6		(a) Nitrogen has less affinity towards halogens(b) Nitrogen halides are covalent	
209.	(c) P_2H_5 Sides of match box have (a) Potassium chlorate,	_	219.	(c) Nitrogen undergoes d^2sp^3 hybridization (d) Nitrogen does not have d -orbitals In the nitrogen family the $H-M-H$ bond angle in the hydrides MH_3 gradually becomes closer to	
	(b) Potassium chlorate, antimony sulphide(c) Antimony sulphide, red phosphorus(d) Antimony sulphide, red lead			90° on going from N to Sb . This shows that gradually [MP PET/PMT 1998; MP PMT 2000]	
210.		compound is tribasic acid [AIIMS 2002] (b) H_3PO_3	bond	(a) The basic strength of hydrides increases (b) Almost pure p -orbitals are used for $M-H$	
211.	(c) H_3PO_4 Which of the following	(d) $H_4P_2O_7$ is manufactured from the	20110	(c) The bond energies of $M-H$ bond increase (d) The bond pairs of electrons become nearer to the central atom	
	molecular nitrogen by b (a) NO_3	acteria [MP PET 1999] (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 (a) B (b) Al	

				s and p-Bloo	ck Elements 785
	(c) N	(d) P		(a) N	(b) <i>P</i>
221.		wing tendencies remains		(c) As	(d) <i>Sb</i>
	unchanged on going down in the nitrogen family (Group-VA)? [MP PMT 1997]		233.		ne following hydrides follow
				the order of	[DPMT 2004]
	(a) Highest oxidation s	tate		(a) $NH_3 > AsH_3 > PH_3 >$	
	(b) Non-metallic character				-
	(c) Stability of hydrides			(b) $SbH_3 > AsH_3 > PH_3 >$	-
222.	 (d) Physical state Which of the following oxy acids of phosphorus is a reducing agent and monobasic (a) H₃PO₂ (b) H₃PO₃ 			(c) $SbH_3 > NH_3 > AsH_3 > ShH_3 > S$	-
			(d) $NH_3 > PH_3 > AsH_3 > SbH_3$ [DCE 2004]		
				Which of the following	
	(c) H_3PO_4	(d) $H_4 P_2 O_6$			[MP PMT 2004]
223.	Bone black is a polymor			(a) PCl_3	(b) BiCl ₃
	(a) Phosphorus	(b) Sulphur		(c) $SbCl_3$	(d) <i>CCl</i> ₄
	(c) Carbon	(d) Nitrogen	235.		l process, the compound
224.	Nitrous oxide is known as [AFMC 2004]			= -	n calcium phosphate is [KCET
	(a) Breathing gas	(b) Laughing gas		(a) Calcium	(b) Phosphine
	(c) exercising gas	(d) Laboratory gas	nont	(c) Phosphorus oxide	(d) Phosphorus
225.	When lead nitrate is he	eated, it gives [MH CET 2003]	-		lowing compound show
	(a) NO_2	(b) <i>NO</i>	230.	sublimation	lowing compound snow
	(c) N_2O_5	(d) N_2O			[AFMC 1995; Pb. CET 2000]
226.	Which element exist	as a solid at 25°C and 1		(a) NH_4Cl	(b) <i>CaCO</i> ₃
	atmospheric pressure a	eric pressure among the following [DCE 200		(c) BaSO ₄	(d) <i>CaHPO</i> ₃
	(a) Br	(b) <i>Cl</i>	227	Number of $P-O$ bonds	3
	(c) Hg	(d) <i>P</i>	23/•		
227.		$O_3 + P_4 O_{10} \to 4HPO_3 + x$, the		(a) 17 (c) 15	(b) 16 (d) 6
	product x is	[MH CET 2003; DPMT 2004]	228	Most acidic oxide is	(u) 0 [Pb. CET 2004]
	(a) N_2O_3	(b) N_2O_5	230.	(a) Na_2O	(b) ZnO
				-	
0	(c) <i>NO</i> ₂	(d) H_2O		(c) MgO	(d) P_2O_5
228.		ogen atom (s) attached to pophosphorous acid is [AIEEE		Which of the following	
	(a) Zero	(b) Two	2005]	(a) NO	[Pb. CET 2003]
	(c) One	(d) Three		(a) <i>NO</i>	(b) NO ₂
229.	• •	is obtained on reacting		(c) N_2O_5	(d) N_2O
	equimolar amounts of two gases at $-30^{\circ}C$ [IIT-JEE (Screeping)xidation number of As in $H_2AsO_4^-$ 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.	allotropic form of phosphorus [IIT-JEE (Screening) 2005]		241.	When plants and am	nimals decay, the organic
			05]	nitrogen is converted into inorganic nitrogen. The	
	(a) Red	(b) White		inorganic nitrogen is ir	
	(c) Black	(d) Yellow		(a) Ammonia(c) Nitrates	(b) Elements of nitrogen
231.	. =	g liberates a gas. The same	_	(C) Nitrates	(d) Nitrides
	gas will be obtained by [IIT JEE (Screening) 2004; BVP:		2004]	Oxygen family	
	(a) Heating NH_4NO_2			Oxygen	ramily
	(b) Heating NH_4NO_3			Mich clare and in farm	lin franctata
	(c) Treating H_2O_2 with $NaNO_2$		1.	Which element is found	n free state PMT 1972, 81, 91; DPMT 1986]
	(d) Treating Mg_3N_2 with	th H_2O		(a) Iodine	(b) Sulphur
232.	The element which for states $+I$ to $+V$ is	rms oxides in all oxidation		(c) Phosphorus	(d) Magnesium

	<u> </u>					
2.	Which of the elementallotropic forms	ts listed below occurs in		(a) H_2O	(b) H_2S	
	anotropic forms	[CPMT 1972]		(c) H_2Se	(d) H_2Te	
	(a) Iodine		16.	Oxygen was discovere	d by [BHU 1987]	
		(b) Copper		(a) Priestley	(b) Boyle	
	(c) Sulphur	(d) Silver		(c) Scheele	(d) Cavandish	
3.	Which forms new compound in air [AFMC 1987]		17.		gives off oxygen on moderate	
	(a) H_2O in air	(b) O_2 in air		heating is	D DIVE 4004. MADE Diber 400-1	
	(c) N_2 in air	(d) Phosphorus in air		(a) Cupric oxide	P PMT 1991; MADT Bihar 1995]	
4.	Which of the following after burning at room			(b) Mercuric oxide		
	temperature gives gaseous oxide [NCERT 1973; CPMT 1981; DPMT 1982; JIPMER 2001]			(c) Zinc oxide(d) Aluminium oxide		
	(a) <i>H</i>	(b) <i>Na</i>	18.		otain oxygen from air by	
	(c) S	(d) <i>He</i>	10.	fractional distillation		
5.	Sulphur molecule is converted into sulphur ion,			(a) Oxygen is in a different group of the peri		
	when it			table from nitroge		
	[DPMT 1980]			(b) Oxygen is more reactive than nitrogen		
	(a) Gains two electrons (b) Loses two electrons				(c) Oxygen has higher b.p. than nitrogen(d) Oxygen has a lower density than nitrogen	
	_	(d) Shares two electrons	19.		air so it is collected over	
6.		berates oxygen gas from	19.	onjgen io denoer man	[CPMT 1980; MP PET 1999]	
	water is			(a) H_2O	(b) Ethanol	
	(a) <i>P</i>	[MP PMT 1993] (b) Na		(c) Mercury	(d) Kerosene oil	
	(c) F	(d) <i>I</i>	20.			
7.	The highest catenation ability is shown by			[CPMT 1991, 99, 2002; AIIMS 1983; BHU 198		
, -		33; MP PET 1993; CPMT 1997]			MP PMT 1985, 92; AFMC 2004]	
	(a) Oxygen	(b) Sulphur		(a) Paramagnetism		
	(c) Selenium	(d) Tellurium			(d) Ferrimagnetism	
8.	Ozone belongs to which group of the periodic table		21.	When oxygen is pas Na_2SO_3 we get	ssed through a solution of	
	(a) V group	(b) VI group		(a) Na_2SO_4	(b) Na ₂ S	
	(c) VII group	(d) None of these		(c) NaHSO 4	(d) <i>NaH</i>	
9.	The number of unpaired electrons in the <i>p</i> -subshell of oxygen atom is		22. [Oxygen does not react Kurukshetra CEE 1991] (a) <i>T</i>	(b) <i>Na</i> [CBSE PMT 1989]	
	(a) 1	(b) 2		(a) I (c) S	(d) Cl	
	(c) 3	(d) 4	23.			
10.		on earth is[MP PET/PMT 1988	3]		ll MEE 1995; RPET 1999, 2000]	
	(a) <i>O</i>	(b) S			ygen (b)Compound of oxygei	
	(c) Se	(d) <i>Te</i>		(c) Isotope of oxygen		
11.	Which of the following	is most electronegative[BHU:	1978] 24.	Ozone is obtained from		
	(a) <i>U</i>	(b) s		(a) By oxidation at hig		
12.	(c) Te	(d) Se		(b) By oxidation using	g a catalyst	
14.	The ability of a substance to assume two or more crystalline structures is called			(c) By silent electric d	lischarge	
	(a) Isomerism	(b) Polymorphism		(d) By conversion at h	iigh pressure	
	(c) Isomorphism	(d) Amorphism	25.		ng statement is true about	
13.	SO_2 is obtained when	[Roorkee 1995]		ozone layer	[NCEPH 4000]	
	(a) Oxygen reacts with dilute sulphuric acid			(a) It is harmful beca	[NCERT 1980] ause ozone is dangerous to	
	(b) Hydrolysis of dilute H_2SO_4			living organism	ause ozone is uangerous to	
	(c) Concentrated H_2SO_4 reacts with Na_2SO_3				cause oxidation reaction can the presence of ozone	
	(d) All of these			=	ecause ozone cuts out the	
14.		hism[BHU 1982; MP PMT 1985]		ultraviolet radiation		
	(a) <i>O</i> (c) <i>Se</i>	(b) S(d) All the above				
15.	Bond angle is minimum					
13.	Dona angie is illillillillilli	[DFM1 1990]				



(d) It is harmful because ozone cuts out the A solution of sulphur dioxide in water reacts with 36. H_2S precipitating sulphur. Here sulphur dioxide important radiation of the sun which are vital for photosynthesis acts as Identify the incorrect statement with respect to [NCERT 1980; MP PMT 1994] (a) An oxidising agent (b) A reducing agent [AIIMS 1992] (c) An acid (d) A catalyst (a) Ozone is formed in the upper atmosphere by a When SO_2 is passed through acidified $K_2Cr_2O_7$ photochemical reaction involving dioxygen solution (b) Ozone is more reactive than oxygen [CPMT 1989, 94] (c) Ozone is diamagnetic whereas dioxygen is (a) The solution turns blue paramagnetic (b) The solution is decolourised (d) Ozone protects the earth's inhabitants by (c) SO_2 is reduced absorbing γ radiations (d) Green $Cr_2(SO_4)_3$ is formed Which one of the following property is not correct 27. for ozone **38.** When SO_2 is passed through cupric chloride [CPMT 1984] solution (a) It oxidises lead sulphide [CPMT 1979, 81, 89, 94] (b) It oxidises potassium iodide (a) A white precipitate is obtained (c) It oxidises mercury (b) The solution becomes colourless (d) It cannot act as bleaching agent (c) The solution becomes colourless and a white 28. Ozone with KI solution produces [CPMT 1987] precipitate of Cu_2Cl_2 is obtained (a) Cl_2 (d) No visible change takes place (c) HI 39. Which of the following is oxidised by SO 2 [BHU 1986] (d) IO₃ The gases respectively absorbed by alkaline (b) $K_2Cr_2O_7$ (a) *Mg* pyrogallol and oil of cinnamon is [CBSE PMT 1989] (d) All of these (c) $KMnO_4$ (a) O_{3} CH_{4} (b) O_2 , O_3 Bleaching action of SO_2 is due to [CPMT 1971, 79, 86] (c) SO_2 , CH_4 (d) N_2O , O_3 (a) Reduction (b) Oxidation Ozone turns trimethyl paper [CPMT 1989] (c) Hydrolysis (d) Its acidic nature (a) Green (b) Violet **41.** A salt of sulphurous acid is called [NCERT 1978] (c) Red (d) Black (a) Sulphate (b) Sulphurate No. of atoms in one molecule of sulphur is (c) Sulphite (d) Sulphide [AFMC 1987, 91; AMU 1985] **42.** Which of the following is acidic (a) 8 (b) 4 [AFMC 1990; JIPMER 1997] (c) 3 (d) None of these (a) SO_3 (b) N_2O **32.** When H_2S is passed through acidified $KMnO_4$, (c) BeO (d) *HgO* we get The final acid obtained during the manufacture of [CPMT 1979, 91; MP PMT 1987] H_2SO_4 by contact process is (a) K_2SO_3 (b) MnO_2 (a) H_2SO_4 (conc.) (b) H_2SO_4 (dil.) (d) Sulphur (c) KHSO₃ (c) H_2SO_4 (d) $H_2S_2O_7$ Copper turnings when heated with concentrated sulphuric acid will give[AFMC 1987; BHU 1999; CBSE PM4200 bout H_2SO_4 which is incorrect (a) Reducing agent (b) Dehydrating agent (a) SO_2 (b) SO₃ (c) Sulphonating agent (d) Highly viscous (c) H_2S (d) O_2 In the reaction Which of the following is used to absorb sulphur $2Ag + 2H_2SO_4 \rightarrow Ag_2SO_4 + 2H_2O + SO_2$ dioxide H_2SO_4 acts as [CPMT 1981] [EAMCET 1980] (a) Reducing agent (b) Oxidising agent (a) Conc. H_2SO_4 (b) KOH solution (c) Catalytic agent (d) Dehydrating agent (c) Water (d) Anhydrous CaCl, 46. In the reaction 35. Which compound acts as an oxidising as well as $HCOOH \xrightarrow{H_2SO_4} CO + H_2O$; H_2SO_4 acts as reducing agent [IIT 1991]





(a) Dehydrating agent (b) Oxidising agent

(d) All of these

(c) Reducing agent

(a) SO₂

(c) Al_2O_3

(b) MnO_2

(d) CrO_3

47. When conc. H_2SO_4 comes in contact with sugar, it becomes black due to

[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) $S_2 O_5^{2-}$
- (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) CdSO₄
- (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

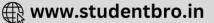
- (b) Convert the metallic silver to silver salt
- (c) Remove undecomposed silver bromide as a soluble complex
- (d) Remove reduced silver
- 59. Hypo is used in photography because of its

[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) CO₂
- (c) *CaO*
- (d) ZnO
- **62.** 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*₅ 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) COCl₂
- (c) Mixture of CO and CO2
- (d) Mixture of O_2 and CO_2
- **71.** Which of the following dissociates to give H^+ most easily

[MP PET 1994]

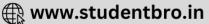
- (a) H_2O
- (b) H_2S
- (c) H_2Te
- (d) H_2Se
- **72.** Superphosphate is the mixture of **[CPMT 1993]**
 - (a) Calcium phosphate and dil. H_2SO_4
 - (b) Sodium phosphate and dil. H_2SO_4
 - (c) Potassium phosphate and dil. H_2SO_4
 - (d) None of these
- **73.** Among KO_2 , NO_2^- , BaO_2 and NO_2^+ 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_2
- (d) CO₂
- **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







76.	In the preparation of su	lphuric acid, V_2O_5 is used		(c) Se	(d) <i>Na</i>
	in the reaction, which is	[CBSE PMT 2001; AFMC 2001]	88.	Which of the following i	is a suboxide [DPMT 2001]
	(a) $S + O_2 \rightarrow SO_2$			(a) Ba_2O	(b) <i>Pb</i> ₂ <i>O</i>
	(b) $2SO_2 + O_2 \rightarrow 2SO_3$			(c) C_3O_2	(d) <i>ZnO</i>
	(c) $SO_2 + H_2O \rightarrow H_2SO_4$		89.	In the manufacture of	sulphuric acid by contact
	(d) $N_2 + 3H_2 \rightarrow 2NH_3$			process, Tyndall box is t	used to [KCET 2003]
77•	Which of the following boiling point	hydrides has the lowest		(a) Filter dust particles(b) Remove impurities	
		[MP PET 1997]		(c) Convert SO_2 to SO_3	
	(a) H_2O	(b) H_2S		(d) Test the presence of	dust particles
	(c) H_2Se	(d) H_2Te	90.	Permono sulphuric acid	is known as [Bihar CEE 1995]
78.		manufacture of H_2SO_4 by		(a) Marshell's acid	(b) Caro's acid
	contact process is	[UPSEAT 1999]		(c) Sulphuric acid	(d) None of these
	(a) Al_2O_3	(b) Cr_2O_3	91.	$KO_2 + CO_2 \rightarrow ? $ (gas)	[CPMT 1997]
	(c) V_2O_5	(d) MnO_2		(a) H_2	(b) N_2
79.	The molecular formula of	-		(c) O_2	(d) <i>CO</i>
	(a) S	MT 1996; MP PET/PMT 1998] (b) S_2	92.	H_2SO_4 acts as dehydra	ating agent in its reaction
		2		with	
0.0	(c) S_4	(d) S_8			[JIPMER 2001]
80.	descicator to dry substa	is not suitable for use in a	Γ/	(a) $H_2C_2O_4$ AIIMS 1996]	(b) <i>Ba(OH)</i> ₂
	(a) Conc. H_2SO_4	(b) Na_2SO_4	L	(c) <i>KOH</i>	(d) Zn
			93.		group shows the highest
_	(c) CaCl ₂	(d) $P_4 O_{10}$		boiling point	[NED DEED as as]
81.	Which shows polymorph			(a) H_2O	[MP PET 2002] (b) H_2S
	(a) <i>O</i> (c) <i>Se</i>	(b) <i>S</i> (d) All		_	_
82.		en family are [MP PET/PMT 19	180	(c) H_2Se	(d) H_2Te
	(a) Non-metals	(b) Metalloids	94.	In presence of moisture	, SO_2 can [BVP 2003]
	(c) Radioactive	(d) Polymorphic		(a) Act as oxidant	(b) Lose electron
83.	The triatomic species	of elemental oxygen is		(c) Gain electron	(d) Not act as reductant
	known as		95.	A gas that cannot be col	
	(a) Agono	[Kerala (Med.) 2002]		(a) N_2	[Kurukshetra CEE 1998] (b) O_2
	(a) Azone(c) Triozone	(b) Polyzone(d) Ozone		-	_
84.		ed through nitric acid, the		(c) <i>SO</i> ₂	(d) <i>PH</i> ₃
-1.	product is	[Kerala (Engg.) 2002]	96.	water on sodium peroxi	is formed by the action of de
	(a) Rhombic S	(b) Prismatic S		(a) H_2	(b) N_2
	(c) Amorphous S	(d) Monoclinic S		(c) O_2	(d) <i>CO</i> ₂
	(e) None of these	(4) 1101100111100	97.	Sulphur on boiling with	NaOH solution gives
85.	Shape of O_2F_2 is similar	to that of [AIIMS 2004]			[Roorkee 1999]
	(a) C_2F_2	(b) H_2O_2		(a) $Na_2S_2O_3 + NaHSO_3$	(b) $Na_2S_2O_3 + Na_2S$
		(d) C_2H_2		(c) $Na_2SO_3 + H_2S$	(d) $Na_2SO_3 + SO_2$
0.6			98.	Quartz is a crystalline v	ariety of
86.	energy	g bonds has the highest			CET 2002; Pb. PMT 2000, 04]
	OJ	[CBSE PMT 1996]		(a) Silicon carbide	(b) Sodium silicate
	(a) <i>Se</i> – <i>Se</i>	(b) <i>Te – Te</i>		(c) Silica	(d) Silicon
	(c) $S-S$	(d) O-O	99.	_	ent for the absorption of
0-	7471-1-1 C +1 C-1111			SO_3 is	



(a) $80\% H_2 SO_4$



[BHU 2004; DPMT 2004]

(b) $98\% H_2SO_4$

(a) O

87. Which of the following is not a chalcogen[CPMT 1999]

(b) S

- (c) $50\% H_2SO_A$
- (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
- **101.** The smog is essentially caused by the presence of [AIEEE 2004]
 - (a) Oxides of sulphur and nitrogen
 - (b) O_2 and N_2
 - (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
- (c) Basic property
- (d) Reducing property
- 103. Oxygen is not evolved on reaction of ozone with
 - [Pb. PMT 2004]

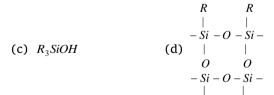
- (a) H_2O_2
- (b) SO,
- (c) Ha
- (d) KI
- **104.** When PbO_2 reacts with conc. HNO_3 the gas evolved is
 - [IIT-JEE (Screening) 2005]
 - (a) NO_2
- (b) O_2
- (c) N_2
- (d) N_2O
- 105. Sulphur in +3 oxidation state is present in [DCE 2003] 6.
 - (a) Sulphurous acid
- (b) Pyrosulphuric acid
- (c) Dithionous acid
- (d) Thiosulphuric acid
- **106.** $SO_2 + H_2S \rightarrow \text{product.}$ the final product is

[Orissa JEE 2005]

- (a) $H_2O + S$
- (b) H_2SO_4
- (c) H_2SO_3
- (d) $H_2S_2O_3$
- **107.** Oncontrolled hydrolysis and condensation, R_3SiCl vields

[Orissa JEE 2005]

- (a) $R_3Si O SiR_3$
- (b) $\{R_3Si O SiR_3\}$



- 108. Ozone deplict due to the formation of following compound in Antarctica [Kerala CET 2005]
 - (a) Acrolein
- (b) Peroxy acetyl nitrate
- (c) SO_2 and SO_3
- (d) Chlorine nitrate
- (e) Formaldehyde

Halogen family

- 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 < HBr
- Phosgene is the common name of 2.

[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 3. 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)
- Bromine is liberated when an aqueous solution of
- 5. 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]

[BHU 1988]

- (a) *HI*
- (b) HBr
- (c) HCl
- (d) HF
- Chlorine was discovered by (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] 9. (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 10.

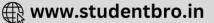
[CPMT 1985]

- (a) Caliche
- (b) Carnellite
- (c) Common salt
- (d) Cryolite
- 11. Which one of the halogen acids is a liquid[MP PMT 1985] (b) HCl (a) *HF*

- (c) HBr
- (d) HI
- 12. Which one of the following acids is the weakest [MP PMT 1985]
 - (a) HClO
- (b) HBr
- (c) HClO₃
- (d) HCl







13.		sodium hydroxide under		(d) Bromine is more	e reactive than	chlorine
	various condition to gi		25. []	N CMME1191X3∃ r is treat	ted with conce	ntrated H_2SO_4
	(a) Sodium chloride	(b) Sodium hypochlorite		redish brown gas e	volved, gas is	[Pb. CET 2003]
	(c) Sodium chlorate	(d) All of these		(a) Mixture of brom	nine and <i>HBr</i>	
14.	Br_2 gas turns starch io	dide paper		(b) HBr		
	[CPMT	1987; AFMC 1987; AMU 1999]		(c) Bromine		
	(a) Blue	(b) Red		(d) None of these		
	(c) Colourless	(d) Yellow	26.	Sea weed is employ	red as a source	of manufacture
15.	When Cl_2 gas is pass	sed through hot and conc.		of		
	solution of KOH, follo	wing compound is formed				2; MP PET 2002]
		[CPMT 1971, 79; BVP 2003]		(a) <i>F</i>	(b) <i>I</i>	
	(a) KCl	(b) KClO ₃		(c) <i>Br</i>	(d) <i>Cl</i>	
	(c) KClO ₂	(d) KClO ₄	27.	Which of the follow	ring is most acti	_
16.	Deacon's process is use	ed in the manufacture of		(-) Cl	(h) n	[MP PET 1990]
	1	[BHU 1979]		(a) Cl_2	(b) <i>Br</i> ₂	
	(a) Bleaching powder	(b) Sulphuric acid		(c) I_2	(d) F_2	
	(c) Nitric acid	(d) Chlorine	28.	Which of the	following re	presents clear
17.	Which of the following	is the weakest acid		electropositive prope		
		IT 1988, 2000; MP PMT 1995;				8; MP PMT 1991]
	MP PET 198	9, 90; Kurukshetra CEE 1998;		(a) <i>F</i>	(b) <i>Cl</i>	
	(a) HE	Roorkee 2000; RPMT 2000]		(c) <i>Br</i>	(d) <i>I</i>	
	(a) <i>HF</i> (c) <i>HBr</i>	(b) <i>HCl</i>	29.	Which statement is		[MP PET 1991]
18.	Which is the most vola	(d) <i>HI</i>		(a) Electronegativit		
10.		979; AIIMS 1980; DPMT 1982;		(b) Electron affinity		
		J & K CET 2005; DPMT 2002]		(c) Melting point of(d) Boiling point of		
	(a) HF	(b) HCl	30.	Strongest reducing		[MP PMT 1990]
	(c) HBr	(d) <i>HI</i>	30.	(a) F^-	_	[MII 1 MII 1990]
19.	On boiling an aqueou	s solution of KClO ₃ with			(b) Cl ⁻	
	iodine, the following p	roduct is obtained[NCERT 1980	0]	(c) Br	(d) <i>I</i> -	
	(a) KIO_3	(b) KCIO ₄	31.	Which of the fol shell electronic con		
	(c) KIO_4	(d) KCl			-	5; MP PMT 1996]
20.	·	lution is disappeared by		(a) $s^2 p^3$	(b) $s^2 p^6$,, <u>.</u>
20.	shaking it with aqueou	s solution of [CPMT 1979, 81; M	P PET	'PMT 1988:		
	МР	PMT 1986; RPMT 1997, 2002]		-	(d) $s^2 p^5$	
	(a) H_2SO_4	(b) <i>Na</i> ₂ <i>S</i>	32.	Chlorine can remov		[MP PET 1990]
	(c) $Na_2S_2O_3$	(d) Na_2SO_4		(a) Br from NaBr		
21.		cl_2 gas may be made by		(b) F from NaF so(c) Cl from NaCl s		
		MnO_4 with a concentrated		(d) F from CaF_2 so		
	solution of	with a concentrated	22	Hydrolysis of which		uring door not
	Solution of	[CPMT 1973]	33.	occur	cii oi the iono	wing does not
	(a) Potassium chloride			occur		[AIIMS 1982]
	(c) Bleaching powder	(d) Hydrochloric acid		(a) <i>VCl</i> ₄	(b) $TiCl_4$	
22.	~ -	ongst the following is[IIT 1989	1	(c) SiCl ₄	(d) CCl ₄	
	(a) $HClO_4$	(b) <i>HClO</i> ₃	34.	Nitric acid converts	•	[MP PMT 1990]
	·	(d) HClO	34.	(a) Iodic acid	(b) Hydro	
	(c) HClO ₂			(c) Iodine nitrate		pentaoxide
23.	Iodine deficiency in die		35.	In KI solution, I_2 in		_
	(a) Nightblindness	(b) Rickets	55.	, -2 -); EAMCET 1992]
_	(c) Goitre	(d) Beri-beri		(a) I^-	(b) KI_2	,, Limen 1992]
24.	Which of the following	is correct [CPMT 1985]			-	
	(a) Iodine is a solid			(c) KI_2^-	(d) KI_3	
	(b) Chlorine is insolubl		36.	Which reaction can	not be used for	
	(c) Iodine is more reac	tive than bromine		of halogen acid		[MP PMT 1989]



- (a) $2KBr + H_2SO_4 \rightarrow K_2SO_4 + 2HBr$
- (b) $NaHSO_4 + NaCl \rightarrow Na_2SO_4 + HCl$
- (c) $NaCl + H_2SO_4 \rightarrow NaHSO_4 + HCl$
- (d) $CaF_2 + H_2SO_4 \rightarrow CaSO_4 + 2HF$
- In dark, which of the following reacts with 37. hydrogen

[MP PMT/PET 1988; Kurukshetra CEE 1998]

- (a) Br_2
- (b) F_2

- (c) I_2
- (d) Cl₂
- 38. The more activeness of fluorine is due to [MP PMT 1990]
 - (a) F-F bond has less energy
 - (b) F_2 is gas at normal temperature
 - (c) Its electronic bond is maximum
 - (d) F-F bond has more energy
- 39. Which of the following after reacting with KI do not remove iodine [MP PET 1989]
 - (a) $CuSO_{4}$
- (b) $K_2Cr_2O_7$
- (c) HNO_3
- (d) HCl
- Aqueous solution of which of the following acids cannot be kept in a bottle of glass [MP PET 1989]
 - (a) *HF*
- (b) *HCl*
- (c) HBr
- (d) HI
- Which of the following pairs is not correctly 41. matched

[MP PET 1993]

- (a) A halogen which is liquid at room temperature-Bromine
- (b) The most electronegative element-Fluorine
- (c) The most reactive halogen-Fluorine
- (d) The strongest oxidizing halogen-Iodine
- Iodine is formed when potassium iodide reacts 42. with a solution of [MNR 1984; MP PET/PMT 1998]
 - (a) $ZnSO_{A}$
- (b) $CuSO_{4}$
- (c) $(NH_4)_2 SO_4$
- (d) Na_2SO_4
- As the atomic number of halogens increases, the 43. halogens

[MP PMT 1991]

- (a) Lose the outermost electrons less readily
- (b) Become lighter in colour
- (c) Become less denser
- (d) Gain electrons less readily
- Which statement is correct about halogens 44.

[EAMCET 1991]

- (a) They are all diatomic and form univalent ions
- (b) They are all capable of exhibiting several oxidation states
- (c) They are all diatomic and form divalent ions
- (d) They can mutually displace each other from the solution of their compounds with metals
- Mark the smallest atom [CPMT 1984, 89] 45.
 - (a) F

(b) Cl

- (c) Br (d) I
- 46. Mark the element which shows only one oxidation state

[BHU 1988; MP PET 2002]

- (a) F
- (b) Cl

(c) Br

47.

- (d) I
- Which of the following arrangement for the three halogens Cl, Br and I when placed in the order of their increasing electron affinity is correct[CPMT 1990]
 - (a) Cl, Br, I
- (b) I, Br, Cl
- (c) Br, Cl, I
- (d) I, Cl, Br
- Which of the following is strongest oxidising agent

[CPMT 1978, 91, 94; MNR 1990; AMU 1983, 84; MP PMT 1991, 92, 96; IIT 1992; UPSEAT 2000]

- (a) F_2
- (b) *Cl*₂
- (c) Br_2
- (d) I_2
- Fluorine is a better oxidising agent than Br_2 . It is 49. due to

[EAMCET 1992]

- (a) Small size of fluorine
- (b) More electron repulsion in fluorine
- (c) More electronegativity of fluorine
- (d) Non-metallic nature of fluorine
- Fluorine is a stronger oxidising agent than chlorine in aqueous solution. This is attributed to many factors except

[Pb. CET 1989]

- (a) Heat of dissociation (b) Electron affinity
- (c) Heat of hydration
 - (d) Ionisation potential
- Mark the element which displaces three halogens 51. from their compounds

[MP PMT 1980, 82; BHU 1984; NCERT 1987]

- (a) F
- (b) Cl
- (d) I
- Which one of the following is the most basic

[CPMT 1975, 77; MP PMT 2001]

(a) I

(b) Br

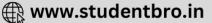
- (c) Cl
- (d) F
- Which of the following will displace the halogen 53. from the solution of the halide [EAMCET 1979]
 - (a) Br_2 added to NaCl solution
 - (b) Cl_2 added to KCl solution
 - (c) KCl added to NaF solution
 - (d) Br_2 added to KI solution
- Fluorine does not form positive oxidation states because

[AIIMS 1987]

- (a) It is most electronegative element
- (b) It forms only anions in ionic compounds
- (c) It cannot form multiple bonding







- (d) It shows non-bonded electron pair repulsion due to small size
- In the isolation of fluorine, a number of 55. difficulties were encountered. Which statement is correct

[NCERT 1983, 86]

- (a) The potential required for the discharge of the fluoride ions is the lowest
- (b) Fluorine reacts with most glass vessels
- (c) Fluorine has great affinity for hydrogen
- (d) Electrolysis of aqueous HF gives ozonised oxygen
- Fluorine reacts with water to give [BHU 1988, 89] 56.
 - (a) HF and O_2
- (b) HF and OF_2
- (c) HF and O_3
- (d) HF, O_2 and O_3
- In which of the following, oxygen has + 2 57. oxidation number [EAMCET 1986]
 - (a) F_2O
- (b) Cl₂O
- (c) Na_2O_2
- (d) Na_2O
- The electrolysis of a certain liquid resulted in the 58. formation of hydrogen at the cathode and chlorine at the anode. The liquid is
 - (a) Pure water
 - (b) H_2SO_4 solution
 - (c) NaCl solution in water
 - (d) CuCl₂ solution in water
- In the preparation of chlorine from HCl, MnO_2 59. acts as

[CPMT 1981]

- (a) Oxidising agent
- (b) Reducing agent
- (c) Catalytic agent
- (d) Dehydrating agent
- Chlorine gas is dried over 60.
- [CPMT 1980]

- (a) *CaO*
- (b) NaOH
- (c) KOH
- (d) Conc. H_2SO_4
- 61. Chlorine can be manufactured from [CPMT 1989]
 - (a) Electrolysis of NaCl
 - (b) Electrolysis of brine
 - (c) Electrolysis of bleaching powder
 - (d) All of these
- 62. When chlorine water is exposed to sunlight, O_2 is liberated. Hence [AFMC 1989]
 - (a) Hydrogen has little affinity to O_2
 - (b) Hydrogen has more affinity to O_2
 - (c) Hydrogen has more affinity to Cl_2
 - (d) It is a reducing agent
- When cold NaOH reacts with Cl2 which of the 63. following is formed [AFMC 1992]
 - (a) NaClO
- (b) NaClO₂
- (c) NaClO₃
- (d) None of these
- Chlorine is used in water for [CBSE PMT 1988] 64.

- (a) Killing germs pollution
- (b) Prevention Ωf
- (c) Cleansing
- (d) Removing dirt
- 65. Chlorine cannot be used
- [MP PET/PMT 1988]
- (a) As bleaching agent
- (b) In sterilisation
- (c) In preparation of antiseptic
- (d) For extraction of silver and copper
- 66. Chlorine acts as a bleaching agent only in presence of

[IIT 1983; DCE 2002]

- (a) Dry air
- (c) Sunlight
- (b) Moisture (d) Pure oxygen
- 67. Euchlorine is a mixture of
- [CPMT 1988]
 - (a) Cl_2 and SO_2
- (b) Cl_2 and ClO_2
- (c) Cl_2 and CO
- (d) None of these
- A gas reacts with CaO, but not with NaHCO₃. 68. The gas is

[AFMC 1987]

- (a) CO₂
- (b) Cl₂
- (c) N_2
- (d) O_2
- When FAMOFTA 939 assed over dry slaked lime at 69. room temperature, the main reaction product is

[CBSE PMT 1992]

- (a) $Ca(ClO_2)_2$
- (b) *CaCl*₂
- (c) CaOCl₂
- (d) $Ca(OCl_2)_2$
- Bromine is obtained commercially from sea water by adding [CPMT 1988]
 - (a) $AgNO_3$ solution
- (b) Crystals of *NaBr*
- (c) Cl₂
- (d) C_2H_4
- In the manufacture of bromine from sea water, the mother liquor containing bromides is treated

[CBSE PMT 1992; MP PMT 2001; BHU 2002; JIPMER 2002]

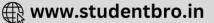
- (a) CO₂
- (b) Cl₂
- (c) I_2
- (d) SO₂
- Br^- is converted into Br_2 by using [CPMT 1987]
 - (a) Cl_2
- (b) Conc. HCl
- (c) HBr
- (d) H_2S
- A salt, which on heating with conc. H_2SO_4 gives violet vapours, is [CPMT 1971]
 - (a) Iodide
- (b) Nitrate
- (c) Sulphate
- (d) Bromide
- When I_2 is dissolved in CCl_4 , the colour that results is

[AFMC 1993]

- (a) Brown
- (b) Violet
- (c) Colourless
- (d) Bluish green
- Which of the following halogen oxides is ionic[CPMT 1989







s and p-Block Elements 793 NCERT 1974: CPMT 1976, 901 (a) ClO₂ (b) BrO_2 (a) HF (b) HCl (d) I_4O_9 (c) I_2O_5 (c) HBr (d) HI KI when heated with conc. H_2SO_4 gives Which of the following hydrogen halides has the 86. [MP PET/PMT 1988] highest boiling point[AIIMS 1980; DPMT 1983; MP PMT 1989 (a) HI (b) I_2 (a) *HF* (b) HCl (c) HBr (d) Hi (c) HIO_2 (d) KIO_3 87. Hydrogen bonding is present in The type of bonding in HCl molecule is[AIIMS 1992] 77. [MP PMT 1989; DPMT 1990; Roorkee 1995] (a) Pure covalent (b) Polar covalent (a) HF (b) HCl (c) Highly covalent (d) H-bonding (c) HBr (d) HI HCl is a gas, but HF is a low boiling liquid. This 78. The type of hybrid orbitals used by chlorine atom is because [EAMCET 1981, 89] in ClO_2^- is (a) H - F bond is strong (b) SP² (a) SP^3 (b) H - F bond is weak (d) None of these (c) SP (c) The molecules aggregate because of hydrogen Which one is the anhydride of HClO₄ bonding in HF [AIIMS 1983; BHU 1983; AMU 1984] (d) HF is a weak acid (a) Cl_2O (b) *ClO*₂ 79. HI cannot be prepared by the action of conc. H_2SO_4 on KI because [MNR 1984] (c) Cl_2O_6 (d) Cl_2O_7 Which of the following halogens is solid at room (a) HI is stronger than H_2SO_4 90. temperature [MP PET 1999; AFMC 1999] (b) HI is more volatile than H_2SO_4 (a) Chlorine (b) Iodine (c) H_2SO_4 is an oxidising agent (c) Bromine (d) Fluorine Which of the following chemical contains chlorine (d) H_2SO_4 forms complex 91. [MP PET 1999] A solution of HCl in water is good conductor 80. (a) Fischer salt (b) Epsom salt while gaseous hydrogen chloride is not. This is (c) Fremy's salt (d) Spirit of salt due to the reason that [NCERT 1980] The element which never acts as reducing agent 92. (a) Water is a good conductor of electricity in a chemical reaction is [Bihar CEE 1995] (b) HCl in water ionises (a) O (b) Li (c) Gas can not conduct electricity but water can (d) C (c) F (d) None of these 93. Concentrated HNO_3 reacts with I_2 to give Sodium chloride when heated with conc. H_2SO_4 [IIT 1989; Roorkee 1995; JIPMER 2001] and solid potassium dichromate gives[CPMT 1981, 84] (a) HI (b) *HOI* (a) Chromic chloride (b) Chromyl chloride (d) HOIO₃ (c) $HOIO_3$ (c) Chromous chloride (d) None of these The formula of some fluorides are given below. 94. Hydrogen bonding does not play any role in 82. Which of them will combine further with fluorine[NCERT boiling of (a) IF_5 (b) NaF [AFMC 1992] (c) CaF_2 (d) SF_5 (a) NH_3 (b) H_2O Which one below is a pseudohalide 95. [AIIMS 1982] (c) HI (d) C_2H_5OH (a) CN^- (b) *ICl* HBr and HI reduce sulphuric acid, HCl can 83. (c) IF_5 (d) $I_3^$ reduce KMnO₄ and HF can reduce[IIT 1981; MP PET 1993] 96. Which one is highest melting halide (a) H_2SO_4 (b) $KMnO_4$ [AIIMS 1982; BHU 1999] (c) $K_2Cr_2O_7$ (d) None of these (a) NaCl (b) NaBr 84. Which has the highest molar heat of vaporisation (c) NaF (d) NaI [CPMT 1991] The above answer is correct because the chosen 97. (a) HF (b) HCl halide has (c) HBr (d) HI [AIIMS 1982] 85. Mark the strongest acid (a) Minimum ionic character

(b) Maximum ionic character

[Bihar MEE 1996; MP PET/PMT 1998;

794 s and p-Block Elements (c) Highest oxidising power (d) Lowest polarity Which of the following oxidizes H_2O to oxygen

(a) Chlorine (b) Fluorine (c) Bromine (d) Iodine

The bleaching action of the bleaching powder is 99. due to the liberation of [MP PMT 1994] (a) Chlorine (b) Molecular oxygen

(c) Nascent oxygen (d) Calcium carbonate

100. Which of the following element is extracted commercially by the electrolysis of an aqueous solution of its compound

[KCET 2002]

[MP PET 1994]

(a) Chlorine (b) Bromine (c) Aluminium (d) Calcium

101. The effective component of bleaching powder is of calcium

[Kerala (Engg.) 2002]

(a) Chlorine (b) Bromine (d) Calcium (c) Aluminium

102. $Na_2S_2O_3 + I_2 \rightarrow \text{Product is}$ [BHU 2003]

(a) Na_2S (b) NaI (c) $Na_2S_4O_6$ (d) S_2

103. Which of the following is prepared by electrolytic method

[CBSE PMT 2001]

(a) Ca (c) S

(b) Sn (d) F_2

104. Beilstein test is used for

[AFMC 1995]

(a) N_2

(b) Cl

(c) Na

(d) CO2

105. Which one will liberate Br_2 from KBr[BVP 2003]

> (a) I_2 (c) HI

(b) SO₂ (d) Cl₂

106. Chlorine dioxide is best prepared by passing dry [Kerala PMT 2003]

(a) Chlorine gas over hot HgO

(b) Chlorine and oxygen gas over hot pt catalyst

(c) Hydrogen chloride and oxygen over silver oxide

(d) Hydrogen chloride over phosphorus pentoxide

(e) Chlorine over hot silver chlorate

107. The mixture of concentrated HCl and HNO_3 made in 3: 1 ratio contains [AIIMS 2003]

(a) ClO₂

(b) NOCl

(c) NCl_3

(d) N_2O_4

108. On exciting Cl_2 molecule by UV light, we get

[UPSEAT 2003]

(a) Cl (c) Cl⁻ (b) Cl⁺

(d) All

109. Which of the following statements is not true

[CBSE PMT 2003]

(a) HF is a stronger acid than HCl

(b) Among halide ions, iodide is the most powerful reducing agent

(c) Fluorine is the only halogen that does not show a variable oxidation state

(d) HOCl is a stronger acid than HOBr

110. The correct order of acidic strength [Pb. CET 2004]

(a) $Cl_2O_7 > SO_2 > P_4O_{10}$

(b) $K_2 > CaO > MgO$

(c) $CO_2 > N_2O_5 > SO_3$

(d) $Na_2O > MgO > Al_2O_3$

111. Halogen acid used in the preparation of aqua regia is

[DPMT 2002]

(a) HBr

(b) HI

(c) HCl

(d) HF

112. NaOCl is used as a bleaching agent and sterilising agent. It can be synthesized by the action of [RPET 2003]

(a) NaCl with H_2O

(b) NH_4Cl with NaOH

(c) Cl_2 with cold and dilute NaOH

(d) Cl2 with hot and concentrated NaOH

113. Metal halide which is insoluble in water is[AIIMS 1996]

(a) AgI

(b) *KBr*

(c) *CaCl*₂

(d) AgF

114. Which one among the following non-metals liquid at 25° C

[MP PMT 1999]

(a) Bromine

(b) Carbon

(c) Phosphorus

(d) Sulphur

115. Bleaching action of chlorine is due to [Bihar CEE 1995]

(a) Oxidation

(b) Reduction

(c) Hydrolysis

(d) Its acidic nature

116. Hydrogen iodide cannot be prepared by the action of conc. H_2SO_4 on potassium iodide because [Bihar CEE 19]

(a) HI is stronger than H_2SO_4

(b) HI is more volatile than H_2SO_4

(c) H_2SO_4 is an oxidising agent

(d) H_2SO_4 forms complex

117. White enamel of our teeth is [Bihar CEE 1995]

(a) $Ca_3(PO_4)_2$

(b) CaF_2

(c) CaCl₂

(d) $CaBr_2$

118. The least active halogen with hydrogen is[DPMT 1996]

(a) Cl

(b) I

(c) Br

(d) F

119. Iodine dissolves readily in

[BHU 1996]

(a) Water

(b) Potassium iodide

(c) Carbon tetrachloride (d) Alcohol





120. Which one of the following compounds in aqueous (b) All halogens form oxy acids solution gives a white precipitate with perchloric (c) All halogens except fluorine form oxy acids acid (d) Only iodine form oxy acids [EAMCET 1997] **131.** When iodine reacts with NaF, NaBr and NaCl (b) KCl (a) NaCl [CPMT 1997] (d) FeCl₃ (c) $MgCl_2$ (a) It gives mixture of F_2 , Cl_2 and Br_2 121. Which of the following sequence is correct with (b) It gives chlorine reference to the oxidation number of iodine[EAMCET 1997] (c) It gives bromine (a) $I_2 > ICl < HI < HIO_4$ (b) $HIO_4 < ICl < I_2 < HI$ (d) None of these (c) $I_2 < HI < ICl < HIO_4$ (d) $HI < I_2 < ICl < HIO_4$ 132. Which is the strongest of the following acids[JIPMER 1999 **122.** The correct order of acidic strength is (a) $HClO_4$ (b) H_2SO_4 [Pb. PMT 1998; AFMC 1998; KCET 2000; Orissa JEE 2005] (c) HCl (d) HNO_3 (a) HF < HCl < HBr < HI133. Hydrogen has a tendency to gain one electron to (b) HCl < HBr < HF < HIacquire helium configuration. In this respect it (c) HBr < HCl < HI < HF(d) HI < HBr < HCl < HF[JIPMER 1999] 123. The chief source of iodine in which it is present (a) Halogens (b) Actinides as sodium iodate is [JIPMER 1997] (c) Transition elements (d) Alkali metals (a) Sea weeds 134. What is the product obtained in the reaction of (b) Caliche $HgCl_2$ and $Hg(CN)_2$ [MP PET 2002] (c) Carnallite (a) $(CN)_2$ (d) Iodine never exists as sodium iodate (b) Addition compound $HgCl_2.Hg(CN)_2$ 124. The lattice energy of the lithium halides is in the following order [Roorkee Qualifying 1998] (c) Hg(CN)Cl(a) LiF > LiCl > LiBr > LiI(d) $Hg[Hg(CN)_2Cl_2]$ (b) LiCl > LiF > LiBr > LiI**135.** The weakest acid HX (X = F, Cl, Br, I) is[BHU 2000] (c) LiBr > LiCl > LiF > LiI(a) *HF* (b) HCl (d) LiI > LiBr > LiCl > LiF(d) HI (c) *HBr* 125. Which has the strong bond [DCE 2001] 136. Bleaching powder is obtained by passing chlorine (a) F - F(b) F - Cl(c) F - Br(d) Cl - B[KCET 2002] (a) Lime stone (b) Quick lime **126.** Iodine and hypo react to produce [DPMT 2001] (c) Slaked lime (d) Pure lime (a) Na_2S (b) Na_2SO_4 **137.** Chlorine is liberated, when we heat [AFMC 1998] (d) Na_2SO_3 (c) $Na_2S_4O_6$ (a) $KMnO_4 + NaCl$ (b) $K_2Cr_2O_7 + MnO_2$ 127. Bleaching powder is correctly represented as[RPMT 1997] (c) $Pb_2(NO_3)_4 + MnO_2$ (d) $K_2Cr_2O_7 + HCl$ (b) CaO (a) CaOCl₂ 138. Which of the following silver compounds finds (c) CaO(Cl) (d) CaCl(OCl) maximum use in photography 128. When chlorine reacts with cold and dilute (a) AgCl (b) AgBr solution of sodium hydroxide, the products (c) AgI (d) $AgNO_3$ obtained are [CBSE PMT 1998] 139. Which of the following halogen does not exhibit (a) $Cl^- + ClO^-$ (b) $Cl^{-} + ClO_{2}^{-}$ positive oxidation state in its compounds (c) $Cl^- + ClO_3^-$ (d) $Cl^{-} + ClO_{4}^{-}$ [EAMCET 1997; AIIMS 2000] (a) Cl (b) Br 129. A one litre flask is full of brown bromine vapour. (c) I (d) F The intensity of brown colour of vapour will not decrease appreciably on adding to the flask some [CBSE1PMT 1998] strength of oxy acids of chlorine follows the order (a) Pieces of marble (b) Carbon disulphide [AIIMS 2000; CBSE PMT 2005] (c) Carbon tetrachloride (d) Animal charcoal





(a) $HClO < HClO_2 < HClO_3 < HClO_4$

(b) $HClO_4 < HClO_3 < HClO_2 < HClO$



130. Which of the following statements is correct[BHU 1997]

(a) Only chlorine and bromine form oxy acids

powder

- (c) $HClO_4 < HClO_3 < HClO < HClO_2$
- (d) None of these
- **141.** Bleaching powder is obtained by treating chlorine with

[Pb. PMT 1999]

- (a) *CaO*
- (b) $CaCO_3$
- (c) CaSO₄
- (d) $Ca(OH)_2$
- **142.** Which statement is not true

[MP PET 2000]

- (a) $Ni(CO)_4$ is diamagnetic
 - (b) BI_3 is a stronger Lewis acid than BF_3
 - (c) Graphite conducts electricity whereas diamond does not
 - (d) CCl_4 is hydrolysed whereas BCl_3 is inert
- **143.** Bleaching powder loses its power on keeping for a long time because **[KCET 2000]**
 - (a) It changes into calcium hypochlorate
 - (b) It changes into calcium chloride and calcium hydroxide
 - (c) It absorbs moisture
- (d) It changes into calcium chloride and calcium chlorate
- **144.** The compound which forms a dative bond with ammonia

[JIPMER 2001]

- (a) *CCl*₄
- (b) BCl₃
- (c) $MgCl_2$
- (d) NaCl
- **145.** The bleaching action of bleaching powder is due to the formation of **[Roorkee 1999]**
 - (a) *CaCl* ,
- (b) CaSO₄
- (c) HClO
- (d) $Ca(ClO_3)_2$
- **146.** Fluorine with dilute *NaOH* gives [MH CET 2000]
 - (a) OF_2
- (b) O_3
- (c) O_2
- (d) HF and O_2
- **147.** Which is not oxidised by MnO_2 [DCE 2003]
 - (a) F

(b) Cl

(c) I_2

- (d) *I*
- **148.** Bromine water reacts with SO_2 to form[AFMC 1995]
 - (a) H_2O and HBr
- (b) H_2SO_4 and HBr
- (c) HBr and S
- (d) S and H_2O
- 149. Which of the following reaction is not feasible

[CBSE PMT PMT 2002]

- (a) $2KI + Br_2 \rightarrow 2KBr + I_2$
- (b) $2H_2O + 2F_2 \rightarrow 2HF + O_2$
- (c) $2KBr + I_2 \rightarrow 2KI + Br_2$
- (d) $2KBr + Cl_2 \rightarrow 2KCl + Br_2$
- **150.** Which of the following has the lowest solubility

[Roorkee 2000]

- (a) CaF_2
- (b) *CaCl*₂

- (c) $CaBr_2$
- (d) CaI₂
- **151.** Which one of the following pairs of substances when mixed, produces chlorine gas at room temperature [IIT 1995]
 - (a) NaCl and MnO_2
 - (b) NaCl and HNO₃ (conc.)
 - (c) NaCl and H_2SO_4 (conc.)
 - (d) HCl (conc.) and $KMnO_4$
- **152.** Concentrated H_2SO_4 cannot be used to prepare HBr from NaBr, because it
 - (a) Reduces HBr
- (b) Oxidises HBr
- (c) Disproportionates *HBr* (d)Reacts slowly with *NaBr* Which of the following halides is least stable and
- **153.** Which of the following halides is least stable and has doubtful existence [IIT 1996]
 - (a) CI_4
- (b) *GeI*₄
- (c) SnI_4
- (d) PbI_4
- **154.** Chlorine cannot displace
- [MP PET 1996]
 - (a) Fluorine from *NaF*
- (b) Iodine from *NaI*
- (c) Bromine from *NaBr* (d) None of these

 55. When fluoride is heated with conc. H SO
- **155.** When fluoride is heated with conc. H_2SO_4 and MnO_2 the gas evolved is **[DPMT 2000]**
 - (a) F_2
- (b) SF
- (c) HF
- (d) None
- **156.** Cl_2 reacts with CS_2 in presence of I_2 catalyst to form

[AFMC 1995]

- (a) $CHCl_3$
- (b) *CCl*₄
- (c) C_2H_5Cl
- (d) C_2H_6
- **157.** Amongst $LiCl, RbCl, BeCl_2$ and $MgCl_2$. Maximum and minimum ionic character will be shown by the compounds

[RPMT 1999]

- (a) LiCl, $MgCl_2$
- (b) RbCl, BeCl₂
- (c) RbCl, MgCl₂
- (d) $MgCl_2$, $BeCl_2$
- **158.** Which is formed when fluorine react with hot and concentrated sodium hydroxide
 - (a) O_2
- (b) O_3
- (c) NaO
- (d) *HF*
- **159.** Which of the following condition is used to find atomic Cl_2 from molecular Cl_2 [CPMT 1996]
 - (a) High temperature, high pressure
 - (b) Low temperature, high pressure
 - (c) High temperature, low pressure
 - (d) Low temperature, low pressure
- **160.** Which one is least basic
- [JIPMER 2000]

- (a) BI_3
- (b) BBr_3 (d) BF_3
- (c) BCl_3
- (u) Br_3
- **161.** On heating $NaCl + K_2Cr_2O_7 + \text{conc.} H_2SO_4$, the gas comes out is [JIPMER 2000]







- (a) O_2
- (b) Cl₂
- (c) CrOCl₂
- (d) CrO_2Cl_2
- **162.** Aqua regia is a mixture of
- [KCET (Med.) 2001]
- (a) $3HCl + 1HNO_3$
- (b) $H_3PO_4 + H_2SO_4$
- (c) $3HNO_3 + 1HCl$
- (d) $HCl + CH_3COOH$
- **163.** Unlike other halogens fluorine does not show higher oxidation states because [MP PET 1997]
 - (a) It is highly electronegative
 - (b) It has no d-orbitals
 - (c) Its atomic radius is very small
- (d) The $\,F^{-}\,$ ion is stable and isoelectronic with neon
- **164.** Which halogen does not show variable oxidation state

[UPSEAT 2003]

- (a) F_2
- (b) Cl₂
- (c) Br₂
- (d) I_2
- **165.** To purify fluorine gas, fumes of HF are removed by

[MH CET 2002]

- (a) Solid NaF
- (b) H_2 gas
- (c) Solid KHF2
- (d) None of these
- **166.** Fluorine is prepared by
 - (a) Oxidation of HF
 - (b) Electrolysis of KF
 - (c) Electrolysis of fused KHF,
 - (d) Decomposition of HgF_2
- **167.** Amongst halogens fluorine is most oxidising because
 - (a) Fluorine has highest electron affinity
 - (b) Fluorine is most electronegative
- (c) Dissociation energy for fluorine molecule is lowest
 - (d) All are correct
- **168.** The alkali metal halides are soluble in water but LiF is insoluble because
 - (a) It is amphoteric
 - (b) The Li-F bond is highly ionic
 - (c) Its lattice energy is high
 - (d) Li⁺ ion is least hydrated
- 169. In which of the following pairs does the first gas bleaches flowers by reduction while the second gas does so by oxidation [Manipal MEE 1995]
 - (a) CO and Cl_2
- (b) SO_2 and Cl_2
- (c) H_2 and Br_2
- (d) NH_3 and SO_2
- 170. Which of the following halogens does not form oxyacid

[MP PET 1997]

- (a) Fluorine
- (b) Chlorine
- (c) Bromine
- (d) Iodine

171. Which of the following molecule is theoritically not possible

[BHU 2002]

- (a) OF_4
- (b) OF_2
- (c) SF_{4}
- (d) O_2F_2
- 172. Iodine is released when potassium iodide reacts with

[UPSEAT 1999]

- (a) $ZnSO_4$
- (b) CuSO 4
- (c) $FeSO_4$
- (d) $(NH_4)_2 SO_4$
- **173.** Which of the following is used in the preparation of chlorine

[CBSE PMT 1999]

- (a) Only MnO₂
- (b) OnlyKMnO₄
- (c) Both MnO_2 and $KMnO_4$
- (d) Either MnO_2 or $KMnO_4$
- **174.** Among Cl^- , Br^- , I^- , the correct order for being oxidise to dihalogen is **[CPMT 1999]**
 - (a) $I^- > Cl^- > Br^-$
- (b) $Cl^- > Br^- > I^-$
- (c) $I^- > Br^- > Cl^-$
- (d) $Br^- > I^- > Cl^-$
- **175.** On heating $KClO_3$, we get
- [CPMT 1999]

- (a) Cl_2O
- (b) *ClO*₂
- (c) ClO_3
- (d) Cl_2O_7
- **176.** For which one of the following properties of halogens the sequence F > Cl > Br > I holds good[MP PET/
 - (a) Electron affinity
- (b) Electronegativity
- (c) Atomic radius
- (d) Boiling point
- 177. Which of the following properties increases on going down from F to I in Group VII-A of the periodic table ?

[MP PMT 1997]

- (a) Electronegativity
- (b) Volatile nature
- (c) Ionic radius
- (d) Oxidising power
- **178.** Among the halogens, the one which is oxidised by nitric acid is

[KCET 2004]

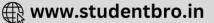
- (a) Fluorine
- (b) Iodine
- (c) Chlorine
- (d) Bromine
- 179. The reaction of the type $2X_2 + S \rightarrow SX_4$ is shown by sulphur when X is [DCE 2003]
 - (a) Fluorine or chlorine
 - (b) Chlorine only
 - (c) Chlorine and bromine only
 - (d) F, Cl, Br all
- **180.** When I_2 is passed through KCl, KF and KBr solutions

[CPMT 2004]

- (a) Cl_2 and Br_2 are evolved
- (b) Cl_2 is evolved







	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) O	(d) None of these	
181.	• •	increases in water in the	5.	Helium was discovere	ed by	
	presence of			(a) Crooks	(b) Rutherford	
	F	[Pb. CET 2002]		(c) Frankland and Lo	ckyer (d)	Dorn
	(a) <i>KI</i>	(b) H_2SO_4	6.	The inert gases are	[СРМТ	1984]
	(c) $KMnO_4$	(d) NH_3		(a) Polyatomic	(b) Triatomic	
102	·	en halides forms salts like		(c) Diatomic	(d) Monoatomic	
102.		llogen atom) [Kerala PMT 2004]	7•	The charcoal maintain		
	(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		
1 Q 2		sodium hydroxide fluorine		(a) Has a saturated of		
103.	reacts to give	[MH CET 2004]		(b) Has one electron i(c) Has eight electron		
	_	(b) $NaF + O_3$		(d) Has two electrons		
	_	(d) $NaF + O_2$	9.	Argon was discovered		1991
404		-	٠,	(a) Rayleigh	(b) Frakland	and
104.	exhibit paramagnetic	ollowing oxides is expected behaviour	Lock			
	(a) CO_2	(b) <i>SO</i> ₂		(c) Jansen	(d) Ramsay	
	-	-	10.	Deep sea divers used	to respirate is a mixtur	e of
_	(c) <i>ClO</i> ₂	(d) SiO ₂		(a) Oxygen and argon		ium
185.	Of the following acids	, the one that is strongest is		(c) Oxygen and nitrog	gen (d) Oxygen	and
	(a) III()	[DPMT 2004]	-	rogen	f	L C
	(a) $HBrO_4$	(b) HOCl	11.	_	forms maximum numl 1976; BHU 1980; CPMT 19	
	(c) HNO_2	(d) H_3PO_3		=	6; MP PMT 2001; Pb.CET	
186.		ig is anhydride of perchloric		(a) <i>Ar</i>	(b) <i>He</i>	
	acid			(c) <i>Xe</i>	(d) <i>Ne</i>	
	(a) Cl O	[CPMT 2004]	12.	Which of the fol	lowing gases exist	more
	(a) Cl_2O_7	(b) Cl_2O_5		abundantly in nature		
	(c) Cl_2O_3	(d) HClO			DPMT 1982,02; CPMT 198	
187.	I_2 dissolves in KI solu	ition due to the formation of		(a) Helium	nipal MEE 1995; MHCET (b) Neon	2003]
		[CPMT 2004]		(c) Argon	(d) Krypton	
	(a) KI_2 and I^-	(b) K^+, I^- and I_2	13.	Which of the followin		
	(c) KI_3^-	(d) None of these	_5.		T 1976, 77; CPMT 1983, 8	6, 90]
	(6)3	(a) None of these		(a) Nitrogen	(b) Fluorine	
	Noble	gases		(c) Neon	(d) Oxygen	
	110010	guood	14.	Nuclear fusion produc	ces	
1.	Which of the fo	llowing outer electronic		(a) Argon	(b) Deuterium	
1.	configuration represe	nts argon[DPMT 1982; CPMT 197 6	5; NCE	R(c) Helium	(d) Krypton	
	0 1	Kurukshetra CEE 1998]	15.	Among the fluorides	below, the one which	does
	(a) ns^2	(b) ns^2np^6		not exist is		
	(c) ns^2np^5	(d) ns^2np^4		() ===	[NCERT 1977; CPMT	1988]
2.	-	sed in isolation of radium		(a) XeF_4	(b) <i>HeF</i> ₄	
۷,	vvincii iiiiiei ai was us	[CPMT 1978, 81, 91]		(c) SF_4	(d) CF_4	
	(a) Lime stone	(b) Pitch blende	16.	The last orbit of argon	n would have electrons	
	(c) Rutile	(d) Haematite			[CPMT 19]	71, 78]
3.	Which is the lightest a			(a) 6	(b) 2	
	(a) Hydrogen	(b) Oxygen		(c) 18	(d) 8	
	(c) Helium	(d) Nitrogen	17.	The electronic configu		
4.	The valency of inert g	ases is		[CPMT 1974,	80, 81; DPMT 1982; MNR	1995]

- (a) $1s^2$, $2s^2 2p^2$
- (b) $1s^2$, $2s^2 2p^6$
- (c) $1s^2$, $2s^2$
- (c) $1s^2$
- **18.** The colour discharge tubes for advertisement mainly contain

[CPMT 1980, 89; MP PET 2002]

- (a) Argon
- (b) Neon
- (c) Helium
- (d) Xenon
- 19. Least chemical activity is shown by [CPMT 1973, 79]
 - (a) Nitrogen
- (b) Argon
- (c) Methane
- (d) Ammonia
- **20.** Noble gases do not react with other elements because

[CPMT 1981]

- (a) They have completely paired up and stable electron shells
- (b) The sizes of their atoms are very small
- (c) Are not found in abundance
- (d) Are monoatomic
- 21. Monazite is source of
 - (a) *He*
- (b) Kr
- (c) Ar
- (d) Ne
- **22.** Which of the following fluorides of Xenon is impossible

[CPMT 1982; Kurukshetra CEE 1998; RPET 1999]

- (a) XeF_6
- (b) XeF_4
- (c) XeF_3
- (d) XeF_2
- **23.** XeF_2 molecule is
 - (a) Square planer
 - (b) Trigonal bipyramidal
 - (c) Trigonal planer
 - (d) Linear
- **24.** XeF_4 on partial hydrolysis produces [AFMC 1995]
 - (a) XeF_2
- (b) $XeOF_2$
- (c) $XeOF_4$
- (d) XeO_3
- **25.** In XeF_2 hybridisation of Xe is
 - (a) sp^2
- (b) sp^3d
- (c) sp^3
- (d) $sp^{3}d^{2}$
- 26. Which one of the following noble gases is the least polarizable[AIIMS 1983; MP PET 1999; Pb. PMT 2001;

JIPMER (Med.) 2002]

- (a) *Xe*
- (b) Ar
- (c) *Ne*
- (d) He
- 27. Which one of the following noble gases is not found in the atmosphere [MP PMT 1993]
 - (a) *Rn*
- (b) Kr
- (c) Ne
- (d) Ar
- **28.** Helium is added to the oxygen supply used by deep sea divers because [MP PMT 1993; MP PET 1997]
- (a) It is less soluble in blood than nitrogen at high pressure $% \left(\frac{1}{2}\right) =\frac{1}{2}\left(\frac{1}{2}\right)$
 - (b) It is lighter than nitrogen

- (c) It is readily miscible with oxygen
- (d) It is less poisonous than nitrogen
- **29.** Which of the following statements is not correct for a noble gas
 - (a) Ar is used in electric bulbs
- (b) Kr is obtained during radioactive disintegration
 - (c) Half life of Rn is only 3.8 days
 - (d) He is used in producing very low temperature
- **30.** Which one of the following configuration represents a noble gas

[CPMT 1976, 83, 89; BHU 1982; Pb. CET 2000 NCERT 1979; IIT Screening 1993; EAMCET 1993]

- (a) $1s^2$, $2s^2 2p^6$, $3s^2$
- (b) $1s^2$, $2s^22p^6$, $3s^1$
- (c) $1s^2$, $2s^2 2p^6$
- (d) $1s^2$, $2s^22p^6$, $3s^23p^6$, $4s^2$
- 31. Which of the following has zero valency[DPMT 1985]
 - (a) Sodium
- (b) Beryllium
- (c) Aluminium
- (d) Krypton
- **32.** The forces acting between noble gas atoms are

[NCERT 1989]

- (a) Vander Waals forces
- (b) Ion-dipole forces
- (c) London dispersion forces
- (d) Magnetic forces
- **33.** Which of the following is the correct sequence of the noble gases in their group in the periodic table

[Manipal MEE 1995]

- (a) Ar, He, Kr, Ne, Rn, Xe(b) He, Ar, Ne, Kr, Xe, Rn
- (c) He, Ne, Ar, Kr, Xe, Rn(d) He, Ne, Kr, Ar, Xe, Rn
- **34.** Which of the following represent nobel gas configuration

[BHU 1995]

- (a) $1s^2$, $2s^22p^6$, $3s^23p^63d^{10}$, $4s^24p^64d^{10}$, $5s^25p^6$
- (b) $1s^2$, $2s^2 2p^6$, $3s^2 3p^6 3d^{10}$, $4s^2 4p^6 4d^{10} 4f^{14}$, $5s^2 5p^6 5d^1$, $6s^2$
- (c) $1s^2$, $2s^22p^6$, $3s^23p^63d^{10}$, $4s^24p^64d^{10}$, $5s^25p^65d^1$, $6s^2$
- (d) $1s^2$, $2s^22p^6$, $3s^23p^63d^{10}$, $4s^24p^64d^{10}$
- **35.** XeF_6 on hydrolysis gives

[MP PET 2000; Pb. PMT 2000; DCE 2002]

- (a) XeO_3
- (b) *XeO*
- (c) *XeO*₂
- (d) *Xe*
- **36.** The correct order of solubility in water for He, Ne, Ar, Kr, Xe, is **[AIIMS 2002]**
 - (a) He > Ne > Ar > Kr > Xe







	800 s and p-Blo	ck Elements	5			
	(b) $Ne > Ar > Kr > He >$ (c) $Xe > Kr > Ar > Ne >$			48.	Which of the following intermolecular forces	ng exhibits the weakest [KCET (Med.) 2001]
	(d) $Ar > Ne > He > Kr >$				(a) He	(b) HCl
37.	In XeF_2 , XeF_4 , XeF_6 the	number of	lone 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	-	10	O	[Roorkee 2000]
	(c) 4, 1, 2	(d) 3, 2, 1			(a) XeF_3	(b) <i>XeF</i> ₄
38.	Noble gases are group				(c) <i>XeF</i> ₅	(d) <i>XeF</i> ₆
	very			50.	Among the following mo	lecule
	() ***		ala (Med.) 2002]	J	(i) XeO_3 (ii) $XeOF_4$ (
	(a) High chemical acti	-				mber of lone pairs on <i>Xe</i>
	(b) Low chemical activ	-		are	Those having same hu	mber of fone pairs on he
	(c) Minimum electron					[AIIMS 2005]
20	(d) Much paramagneti		water [CDMT 2002]		(a) (i) and (ii) only	(b) (i) and (iii) only
39.	Which noble gas is mo (a) He	(b) <i>Ar</i>	water [CPM1 2002]		(c) (ii) and (iii) only	(d) (i),(ii) and (iii)
	(c) Ne	(d) <i>Xe</i>		51.	Who among the followi compound of noble gas	ng first prepared a stable [MP PET 1999]
40.	Gradual addition of el				(a) Rutherford	(b) Rayleigh
	gases causes a decreas		[MP PET 1997]		(c) Ramsay	(d) Neil Bartlett
	(a) Ionization energy(c) Boiling point	(b) Atomi (d) Densi		52.	The last member of iner	t gas elements is[MP PMT 1999]
41.	Which of the following		-		(a) Helium	(b) Neon
41.	octet of electrons in it				(c) Argon	(d) Radon
	(a) Neon	(b) Rador		53.	Which of the following g	·
	(c) Argon	(d) Heliu	m			[CPMT 2000; Pb. CET 2002]
42.	The low chemical read	ctivity of the	rare gases can		(a) Ne	(b) He
	be attributed to their		[Pune CET 1998]		(c) Kr	(d) All of these
	(a) Being non-metals			54.	helium is incorrect	ving statements regarding [AIEEE 2004]
	(b) Having high ioniza	tion energie	S			ice and sustain powerful
	(c) Being gases		•.•		superconducting ma	
	(d) Found in nature in	=			(b) It is used as a cryoge	enic agent for carrying out
43.	Percentage of Ar in ai		[CPMT 1989]		experiments at low t	1
	(a) 1%	(b) 2%				gas balloons instead of
4.4	(c) 3% Which of the following	(d) 4%	nined by direct		inflammable	it is lighter and non-
44.	reaction of constituent		[MP PET 1994]		(d) It is used in gas-cool	ed nuclear reactors
	(a) <i>XeF</i> ₂	(b) <i>XeF</i> ₄		55.	Which of the following i	
	(c) XeO_3	(d) <i>XeF</i> ₆			Č	[Pb. CET 2002]
45.	Fluorine forms chemic	-	ls with[MD DMT 100	41	(a) Kr	(b) He
45.	(a) He	(b) <i>Ne</i>	13 ***1011[1411 1 1411 199	41	(c) Ne	(d) Ar
	(c) Ar	(d) <i>Xe</i>		56.	The oxidation number of	f xenon in $XeOF_2$ is[J & K 2005]
46.	Which of the following		oridisation[DCE 200	11	(a) Zero	(b) 2
-	(a) <i>XeO</i> ₃	(b) <i>BCl</i> ₃	•	-	(c) 4	(d) 3
	-	2		57•		nighest boiling point[BCECE 2005]
	(c) XeF ₄	(d) BBr ₃	1 77 6		(a) <i>Xe</i>	(b) <i>Ar</i>
47.	Which element out of			_	(c) Kr	(d) He
	least number of compo		[MP PMT 1995]	58.	_	s an inert gas [AFMC 2005]
	(a) <i>He</i> (c) <i>Kr</i>	(b) <i>Ar</i> (d) <i>Xe</i>			(a) H_2	(b) O_2
	(C) M	(u) Ae			(c) N_2	(d) Argon
				59.	Which of the following i	s 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 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
- $BaSO_4$ and carbon on heating reacts to produce 5.

[Pb. PMT 2004]

- (a) $Ba + SO_2 + CO_2$
- (b) BaS + CO
- (c) $BaS + O_2 + SO_2$
- (d) $BaCO_3 + S + O_2$
- 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
- 7. Fusion mixture is

[CPMT 2002]

- (a) $Na_2CO_3 + K_2CO_3$
- (b) $Na_2CO_3 + NaHCO_3$
- (c) $Na_2CO_3 + NaOH$
- (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) KO₂
- (4) H_2O_2

Codes:

(a) A В C D

3 4 2 1

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

[KCET 1998]

[NDA 1999]

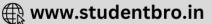
- (a) $98\%H_2SO_4$
- (b) $80\%H_2SO_4$
- (c) 20% oleum
- (d) $90\% H_2 SO_4$
- Mark the oxide which is amphoteric in character 11. [MP PMT 2000]
 - (a) CO_2
- (b) SiO₂
- (c) SnO_{2}
- (d) CaO
- Concentrated aqueous sodium hydroxide can separate mixture of [MNR 1995]

The composition of the common glass is[DCE 2004]

- (a) Al^{3+} and Sn^{2+}
- (b) Al^{3+} and Fe^{3+}
- (c) Al^{3+} and Zn^{2+}
- (d) Zn^{2+} and Pb^{2+}
- (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.	The metal which does	not form ammonium nitrate		(c) Sea shells	(d) Dolomite
	by reaction with dilute (a) <i>Al</i>		25.[]		llowing statements is false for
	(c) <i>Pb</i>	(d) Mg		aikaii illetais	[MNR 1994; MP PET 2001]
5.		pair of electrons in $XeOF_4$ is		(a) Lithium is the	strongest reducing agent
٠.	rotar number of fone p	[IIT-JEE (Screening) 2004]		(b) Na is amphot	
	(a) 0	(b) 1		(c) Li^+ is exception	
	(c) 2	(d) 3		_	als give blue solution in liquid
6.		elationship between the <i>pHs</i>		ammonia	and give dide condition in inquid
		s of sodium oxide (pH_1) ,	26.	Solubility of iodin	ne in water is greatly increased
	sodium sulphide (pH_2)	, sodium selenide (pH_3) and		-	of iodide ions because of the
	sodium telluride (pH_4)			formation of	
		[CBSE PMT 2005]		(a) I	[IIT 1994]
	(a) $pH_1 > pH_2 = pH_3 > p$	bH_4		(a) I_2	(b) I_3
	(b) $pH_1 < pH_2 < pH_3 < p$	bH_4		(c) I_3^-	(d) I^-
	(c) $pH_1 < pH_2 < pH_3 = p$	·	27.	-	water of sulphates down the Be $>> Ca > Sr > Ba$. This is due to
	(d) $pH_1 > pH_2 > pH_3 > p$	bH_4		group is be > mg >	[CBSE PMT 1995]
7.	Which one of the follo	owing is not an amphoteric		(a) High heat of	solvation for smaller ions like
	substance		Be^{2+}	(a) 111811 11040 01	2017-011-101 211-011-01 10110 11110
	()	[KCET 2004]		(b) Increasing mo	lecular weight
	(a) HNO_3	(b) HCO_3^-		(c) Decreasing lat	_
	(c) H_2O	(d) NH_3		(d) Increase in me	
8.	~ -	buffer group of the periodic	28.	Magnesium burns	in air to give
	table	Inh Company			[CPMT 1988, 89; AFMC 1987]
	(a) I	[Pb. CET 2004] (b) VII		(a) <i>MgO</i>	(b) Mg_3N_2
	(c) VIII	(d) Zero		(c) $MgCO_3$	(d) MgO and Mg_3N_2
9.		salt is insoluble in water	both	l	
,		[MP PET 2004]	29.	Philospher's woo	ol when heated with BaO at
	(a) CuSO ₄	(b) $CdSO_4$		$1100~^{o}C$ gives a co	mpound. Identify the compound[
	(c) <i>PbSO</i> ₄	(d) $Bi_2(SO_4)_3$		(a) $BaZnO_2$	(b) $Ba + ZnO_2$
ο.	Which of the following	oxides is the most acidic		(c) BaCdO ₂	(d) $BaO_2 + Zn$
	-	BSE PMT 1999; MP PMT 2002]	30.	Which of the follo	wing oxide is diamagnetic
	(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
1.	Whose bond energy is	maximum		(c) NO_2	(d) N_2O_5
		[CPMT 1988; MP PMT 1990]	31.	-	lowing salt becomes plaster of
	(a) F_2	(b) Cl ₂	J=-		propriately hydrated [CPMT 1985]
	(c) Br ₂	(d) I_2		(a) $ZnCO_3$	(b) CaSO ₄
2.	Calcium cyanide on tr pressure gives <i>NH</i> ₃ ar	reatment with steam under		(c) MgSO ₄	(d) CaCO ₃
	(a) $CaHCO_3$	(b) <i>CaO</i>	32.	The number of e alkaline earth met	lectron and proton in the third all ion will be
	(c) <i>Ca(OH)</i> ₂	(d) CaCO ₃		(a) $\frac{e}{20}, \frac{p}{20}$	(b) $\frac{e}{}$ $\frac{p}{}$
3.	-	n, on complete ozonisation,		20 ' 20	(b) $\frac{e}{18}, \frac{p}{20}$
-	form volumes of o		[]	DPMT <u>20</u> 00 <u>0]</u> 18, 18	(d) $\frac{e}{19}, \frac{p}{20}$
	(a) 4	(p) 3			
	(c) 2	(d) 6	33.		f alkaline earth metals have the
4.	The substance not like	ly to contain $CaCO_3$ is		following magnet	
	[AIEEE 2003]				T 1998; RPMT 2000; JIPMER 2002]
	(a) A marble statue	(b) Calcined gypsum		(a) Diamagnetic	(b) Paramagnetic





- (c) Ferromagnetic
- (d) Diaferromagnetic
- **34.** Which of the following is the life saving mixture for an asthma patient [MP PMT 2001]
 - (a) Mixture of helium and oxygen
 - (b) Mixture of neon and oxygen
 - (c) Mixture of xenon and nitrogen
 - (d) Mixture of argon and oxygen
- 35. Which would quickly absorbs oxygen

[CBSE PMT 1992; MP PET 1995]

- (a) Alkaline solution of pyrogallol
- (b) Conc. H_2SO_4
- (c) Lime water
- (d) Alkaline solution of CuSO 4
- **36.** Nitrogen is liberated by the thermal decomposition of only

[IIT 1991]

- (a) NH_4NO_2
- (b) NaN₃
- (c) $(NH_4)_2 Cr_2 O_7$
- (d) All the three
- **37.** Red phosphorus is less reactive than yellow phosphorus because

[DPMT 1982; JIPMER 1999; CBSE PMT 1999; RPET 2003]

- (a) Its colour is red
- (b) It is highly polymerised
- (c) It is hard
- (d) It is insoluble in C_2H_5OH
- **38.** Carbon differs from other elements of the group. Which is the false statement
 - (a) Due to its marked tendency to form long chains (catenation)
 - (b) Due to its unique ability to form multiple bonds
 - (c) Due to *d*-orbital in penultimate shell
 - (d) Due to its limitation of co-ordination number 4
- **39.** Which of the following oxide does not form acidic aqueous solution [CPMT 2004]
 - (a) N_2O_3
- (b) NO_2
- (c) N_2O_5
- (d) NO
- **40.** Which of the following is in the increasing order of the ionic character [JIPMER 2002]
 - (a) $PbCl_4 < PbCl_2 < CaCl_2 < NaCl$
 - (b) $PbCl_2 < PbCl_4 < CaCl_2 < NaCl$
 - (c) $PbCl_2 < PbCl_4 < NaCl < CaCl_2$
 - (d) $PbCl_4 < PbCl_2 < NaCl < CaCl_2$
- **41.** Silicon chloroform is prepared by [MH CET 1999]
 - (a) Si + HCl
- (b) $SiCl_4 + H_2O$
- (c) $SiF_4 + NaF$
- (d) $H_2SiF_6 + Cl_2$

- **42.** KO_2 (potassium superoxide) is used in oxygen cylinders in space and submarines because it [AIEEE 2002
 - (a) Absorbs CO_2 and increases O_2 content
 - (b) Eliminates moisture
 - (c) Absorbs CO₂
 - (d) Produces ozone
- 43. Fire extinguishers contain H_2SO_4 and [AFMC 1980]
 - (a) $CaCO_3$
- (b) Na_2CO_3
- (c) NaHCO₃
- (d) $NaHCO_3$ and Na_2CO_3
- **44.** Which is insoluble in water
- [CPMT 2003]

- (a) H_2S
- (d) CaF_2
- (c) $Ca(NO_3)_2$
- (--)

(b) $HgCl_2$

- **45.** Which of the following halides is most acidic[KCET 1996]
 - (a) PCl_3
- (b) $SbCl_3$
- (c) $BiCl_3$
- (d) CCl_4
- **46.** The stability of the following alkali metal chlorides follows the order
 - (a) LiCl > KCl > NaCl > CsCl
 - (b) CsCl > KCl > NaCl > LiCl
 - (c) NaCl > KCl > LiCl > CsCl
 - (d) KCl > CsCl > NaCl > LiCl
- **47.** The reaction of $Na_2S_2O_3$ with iodine gives

[CPMT 1971, 80, 81; DPMT 1983, 90; MP PMT 1985; EAMCET 1990; BHU 1980]

(a) Sodium sulphide (b) S

- (b) Sodium sulphite
- (c) Sodium sulphate
- (d) Sodium tetrathionate
- **48.** Which one of the following is the true covalent oxide of iodine [MP PET/PMT 1988]
 - (a) I_2O_4
- (b) I_2O_5
- (c) I_2O_7
- (d) I_2O_9
- **49.** Lithium aluminium hydride acts as **[CPMT 1994]**
 - (a) Oxidising agent
- (b) Reducing agent
- (c) Both the above
- (d) None of these
- **50.** The mixture of conc. *HCl* and potassium chlorate on heating gives [Roorkee 2000]
 - (a) Cl_2 only
- (b) ClO_2 only
- (c) $Cl_2 + ClO_2$
- (d) $Cl_2 + ClO_2 + ClO_3$
- 51. When SO_2 is passed through acidified solution of H_2S

[CPMT 1973, 81, 93]

- (a) H_2SO_4 is formed
- (b) H_2SO_3 is formed
- (c) Sulphur is precipitated (d)
-) None of these
- **52.** Four reactions are given below
 - (i) $2Li + 2H_2O \rightarrow 2LiOH + H_2$
 - (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

- (a) CaCO₃, KHCO₃, NaHCO₃
- [AFMC 1987]
- (b) NaHCO₃, KHCO₃, CaCO₃
- (c) KHCO₃, NaHCO₃, CaCO₃
- (d) CaCO₃, NaHCO₃, KHCO₃
- 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) CaCN₂
- The following acids have been arranged in the 55. 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
- **56.** 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
- H_3PO_2 is the formula for one of the phosphorus 57. 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 AlCl₃
 - (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]
- (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) CO₂
- (d) ClO_2
- Which of the following product is formed when SiF₄ 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
 - (b) $Fe_2O_3 + Al$
 - (a) $Cr_2O_3 + Al_2O_3$ (c) $Fe_2O_3 + Al_2O_3$
- (d) $Al_2O_3 + 2Cr$
- **68.** The colour of liquid O_2 is
- [BVP 2004]

[Pb. CET 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) $O_2 + N_2$
- 70. One mole of magnesium nitride on the reaction 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]

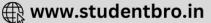
- (a) $CaCO_3 + NH_3$
- (b) $CaHCO_3 + NH_3$
- (c) $CaO + NH_3$
- (d) $Ca(OH)_2 + NH_3$



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

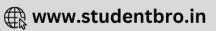






				s a	and p-Block Elements 805
(a) (b)	reason is the	rtion and reason are true and the correct explanation of the assertion. tion and reason are true but reason is		Reason :	The catenated $-O-O-O$ changes are less stable as compared to $O=O$ molecule. [AIIMS 2001]
	not the correc	ct explanation of the assertion.	10.	Assertion:	Potassium and caesium are used in photo-electric cells.
(c)	-	s true but reason is false.		Reason :	Potassium and caesium emit
(d) (e)	-	on and reason both are false. s false but reason is true.		Reason .	electrons on exposure to light.[AIIMS 2002]
(6)	ij assertion is		11.	Assertion :	The fluorine has lower reactivity.
1.	Assertion :	Sulphate is estimated as BaSO ₄ and		Reason :	F-F bond has low bond
1.	Assertion.	•			dissociation energy. [AIIMS 2002]
			12.	Assertion:	Halogens do not occur in free state.
	Reason :	Ionic radius of Mg^{2+} is smaller than		Reason :	Halogens are highly reactive. [AIIMS 1994]
			13.	Assertion:	Lithium forms Lithium oxide (LiO_2) .
2.	Assertion :	Amongst the halogens fluorine can oxidise the elements to highest oxidation state.		Reason :	N_2 molecule have unpaired electrons.
	Reason :	Due to small size of fluoride ion, it			[AIIMS 1995]
			14.	Assertion :	Liquid NH_3 is used for
		fluorine. Hence reverse reaction			refrigeration.
3.	Assertion :	takes place more easily. [IIT 1996] <i>HNO</i> ₃ is a stronger acid than		Reason :	Liquid NH_3 quickly vaporises.
3.	Assertion:	HNO_2 .			[AIIMS 1995]
	Reason :	-	15.	Assertion:	$Al(OH)_3$ is insoluble in NH_4OH but
	Reason :	In HNO_3 there are two nitrogen-to-oxygen bonds whereas in HNO_2			soluble in NaOH.
		thora is only one	_	Reason :	NaOH is strong alkali. [AIIMS 1997]
4.	Assertion :	there is only one. The Value of Vander Waal's	16.	Assertion :	Boron is metalloid.
4.	Assertion:	constant 'a' is larger for ammonia		Reason :	Boron shows metallic nature. [AIIMS 1997]
		than for nitrogen.	17.	Assertion:	Inert gases are monoatomic.
	Reason :	Hydrogen bonding is present in ammonia.		Reason :	Inert gases have stable configuration.
		[IIT 1998]	18.	Assertion :	[AIIMS 1999] Magnesium continue to burn in
5.	Assertion:	Action forms fluorides.	10.	Assertion .	nitric oxide.
	Reason :	Due to the strong electronegativity of fluorine . [AIIMS 2001]		Reason :	During burning heat evolved do not decompose <i>NO</i> . [AIIMS 2001]
6.	Assertion:	Chlorine and sulphur dioxide both are bleaching agents.	19.	Assertion :	Anhydrous BaO_2 is used for
	Reason :	The bleaching action of chlorine	_		preparing H_2O_2 .
		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	-•		benzene is unreactive compound.
		at elevated temperatures (On 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]
			21.	Assertion:	Halogens absorb visible light.
8.	Assertion :	Covalency of oxygen is three.		Reason :	All halogens are coloured.[AIIMS 2002]
	Reason :	quite common but dinegative anion	22.	Assertion :	Barium is not required for normal biological function in human.
		of sulphur (S^{2-}) is less common.[AIIMS 2	001]	Reason :	Barium does not show variable
9.	Assertion:	At room temperature, oxygen exists			oxidation state. [AIIMS 2003]
			23.	Assertion :	The $O-O$ bond length in H_2O_2 is
		exists as solid.			shorter than that of O_2F_2 .
				Reason :	H_2O_2 is an ionic compound.[AIIMS 2003]





Assertion: PbI_{A} is a stable compound. 24.

Reason Iodide stabilizes higher oxidation

state.

[AIIMS 2003]

Assertion: *Mq* is not present in enamel of human 25.

teeth.

Reason Mg is an essential element for

biological functions of human. [AIIMS 2004]

Radium is most abundant s-block 26. Assertion:

element.

s-block elements Reason nonare

radioactive in nature.

LiCl is predominantly a covalent Assertion:

compound.

Reason Electronegativity difference

between *Li* and *Cl* is too small.

28. Assertion: The first ionization energy of Be is

greater than that of B.

2p-orbital is lower in energy than 2s-Reason

orbital.

Assertion: The alkali metals can form ionic 29.

hydrides which contains

hydride ion.

Reason alkali metals have

electronegativity, their hydrides conduct electricity when fused and

liberate hydrogen at the anode.

30. Assertion: does not impart characteristic colour to the bunsen

flame.

Reason Due to its very high ionization

> energy, beryllium requires a large amount of energy for excitation of

the electrons.

Assertion: Potassium is not obtained by the 31.

electrolysis of fused KCl.

Potassium vapourises at the melting Reason

point of KCl.

Assertion: Helium and beryllium have similar 32.

outer electronic configuration of

the type ns^2 .

Both are chemically inert. Reason

Assertion: Na_2SO_4 is soluble while $BaSO_4$ is 33.

insoluble.

Lattice energy of $BaSO_4$ exceeds its Reason

hydration energy.

Alkali metals impart colour to the Assertion: 34.

Reason Their ionisation energies are low.

35. Assertion: Superoxides of alkali metals are

paramagnetic.

Superoxides contain the ion O^{-2} Reason

which has one unpaired electron.

36. Assertion: Although PF_5 , PCl_5 and PBr_3 are

known, the pentahalides of nitrogen

have not been observed.

Reason Phosphorus has lower

electronegativity than nitrogen.

Assertion: The electronic structure of O_3 is 37.

Reason

Structure is not allowed because octet around O cannot be expanded.

38. Assertion: Sulphuric acid is more viscous than

water.

Reason Concentrated sulphuric acid has a

great affinity for water.

PCl₅ is covalent in gaseous and Assertion:

liquid states but ionic in solid state.

PCl₅ in solid state consists of Reason

> tetrahedral cation PCl_{A}^{+} and

octahedral PCl₆ anion.

Assertion: Among nitrogen halides NX_3 , the

dipole moment is highest for

 NI_3 and lowest for NF_3 .

Reason Nitrogen halides NX_3 , have trigonal

pyramidal structure.

Assertion: White phosphorus is stored under 41.

water.

White

Reason

phosphorous is highly reactive and catches fire

spontaneously in air.

Al forms $[AlF_6]^{3-}$ but B does not Assertion: 42.

form $[BF_6]^{3-}$.

Reason B does not react with F_2 .

 NO_3^- is planar while NH_3 Assertion: 43.

pyramidal.

N in NO_3^- is sp^2 hybridized but in Reason

 NH_3 it is sp^3 – hybridized.

Assertion: Si - Si bonds are much stronger

then Si - O bonds.

Silicon forms double bonds with Reason

itself

The 45. Assertion: S-S-S bond angle in

 S_8 molecule is 105°.

 S_8 has a V-shape. Reason



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.

inglier as compared to diprotrain.

53. Assertion: Water can be transformed from

liquid to solid state only.

 $Reason \quad : \quad The \ distribution \ of \ water \ over \ the$

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 : 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

molecule is fixed in the space.

59. Assertion :

Calgon is used for removing Ca^{2+}

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

Reason : Cal

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

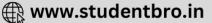
paramagnetic. [AIIMS 2005]



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







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	а	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	а
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	а
21	С	22	С	23	d	24	С	25	а
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	а
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
040				218	d	219	d	220	С
216	b	217	а	210	u	213			·
216	b a	217	a	223	а	224	b	225	a

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

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	а
116	С	117	b	118	b	119	b	120	b
121	d	122	а	123	b	124	а	125	а
126	С	127	d	128	а	129	а	130	С
131	d	132	а	133	а	134	а	135	а
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	а
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	a
166	С	167	С	168	С	169	b	170	а
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	a
186	а	187	С						

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	а
66	С	67	b	68	d	69	а	70	а
71	а								

Noble gases

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	a
11	С	12	b	13	d	14	С	15	b



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

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 - MgCO₃. CaCO₃

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

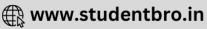
- **14.** (d) Anhydrous form of Na_2CO_3 does not decompose on heating even to redness. It is a amorphous powder called soda ash.
- 17. (c) Fehling's solution is a mixture of Alk. $CuSO_4 + Na K$ tartarate (Rochelle salt)
- 19. (b) $2K+2HCl \rightarrow 2KCl+H_2$ (violent reaction).
- 20. (b) Although 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 ∞ 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) $2NaCl \xrightarrow{\text{Electroly sis}} 2Na + Cl_2$ Anode
 Anode
- **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.

$$KClO_4 \rightleftharpoons K^+ + ClO_4^-$$
; $H_2O \rightleftharpoons OH^- + H^+$
 $KOH \atop Strong HClO_3$

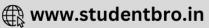
- **91.** (c) Carbon dioxide does not help in burning, also it forms carbonate with alkali metals.
- **92.** (a) When carbonate are heated they decompose to form the oxide. Sodium carbonate and potassium carbonate do not decompose. The carbonate become more difficult to decompose as we go down the group.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

$$2Al + 2NaOH + 2H_2O \rightarrow \underbrace{2NaAlO_2}_{\text{Sodium meta aluminate}} + 3H_2 \uparrow$$

- **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 $H_2{\cal O}$. 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) $Na_2SO_3 + S \xrightarrow{NaOH} Na_2S_2O_3$ Sod. thiosulph at
- **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) $2NaCl \xrightarrow{\text{Electric current}} 2Na^+ + 2Cl^-$ 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 $CaO + H_2O \rightarrow CaOH_2$ burns to form sodium oxide and sodium peroxide.
- **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) CaCl₂ because it is hygroscopic
- **7.** (d) Setting of plaster of paris is exothermic process

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

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) $Anode 2Cl^- \rightarrow 2Cl + 2e^-, Cl + Cl \rightarrow Cl_2$ 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 Dehydration 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

 Sleaching

 Rowyder
- **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 P
 Atomic radii 1.60 1.43 1.32 1.28
 (Å) as we move across the period nuclear charge increases, hence, size decreases.
- $\frac{\textit{MgCl}_2.5 \textit{MgO.xH}_2\textit{O}}{\text{Magnesia cement or sorrel cement}}$ **31.** (d) $\textit{ZnS} + \textit{BaSO}_4$ is lithopone used as white

(b) $MgCl_2.6H_2 + 5MgO + xH_2O \rightarrow$

30.

- pigment.

 26 (d) Aqueous CaCl or hydrated CaCl can
- **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.
- **64.** (a) (i) Small atomic size.
 (ii) High electronegativity
 (iii) Absence of *d* orbitals
- **65.** (a) $Ba(OH)_2 > Sr(OH)_2 > Ca(OH)_2 > Mg(OH)_2$ Solubility decreasing order.
- **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. $Cl_2 + H_2O \rightarrow HCl + HClO$ $\frac{HClO \rightarrow HCl + O}{Cl_2 + H_2O \rightarrow 2HCl + O}$

Coloured matter + nascent oxygen - colourless matter

- **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.

$$\textit{CaNCN} + 2H_2O \rightarrow \textit{CaCO}_3 + \textit{NH}_2 \\ \textit{CONH}_2 \\ \textit{Urea}$$

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

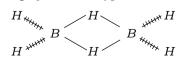
- **104.** (b) Due to electropositive and reactive in nature, magnesium is readily converted into positive ions on contact with iron pipes and hence, iron pipes remains as it is.
- 105. (c) A binary compound is one made of two different elements. These can be one of each element such as CuCl or FeO. 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) $MgCl_2 + 2NaHCO_3 \rightarrow Mg(HCO_3)_2(aq) + 2NaCl$ $Mg(HCO_3)_2(aq) \stackrel{\Delta}{\longrightarrow} MgCO_3 \downarrow + H_2O + CO_2 \uparrow$ (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

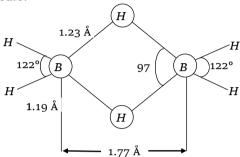


119 pm

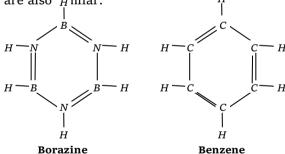
134 pm



- 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.



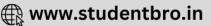
- **15.** (c) $2H_3BO_3 \rightarrow B_2O_3 + 3H_2O$.
- **16.** (a,c,d) Al_2Cl_6 , In_2Cl_6 , Ga_2Cl_6
- **17.** (a) Liquified *Ga* expand on solidification *Ga* is less electropositive in nature, It has the weak metallic bond so it expand on solidification.
- **18.** (d) $Al_2Cl_6 + 12H_2O = 2[Al(H_2O)_6]^{3+} + 6Cl^{-1}$
- **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 illar.



- **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) $Al \rightarrow III \text{ group} \rightarrow \text{Forms } Al_2O_3$
- **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 $Al_2O_3.2H_2O + Na_2CO_3 \rightarrow 2NaAlO_2 + CO_2 + 2H_2O$ $2NaAlO_2 + 3H_2O + CO_2 \xrightarrow{333 K}$ $2Al(OH)_3 \downarrow + Na_2CO_3$ $2Al(OH)_3 \xrightarrow{1473 K} Al_2O_3 + 3H_2O$
- **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.



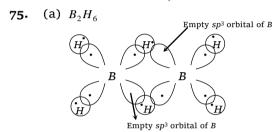


- **67.** (c) In electrolytic method of obtaining aluminium from purified bauxite, cryolite is added to charge because it reduces the melting point of Bauxite (from $1200^{\circ}C$ to $800^{\circ}-900^{\circ}C$) and also it increases electrical conductivity of mixture.
- **68.** (a) Hoop's process \Rightarrow 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. $BF_3 < BCl_3 < BBr_3 < BI_3$.

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 π 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.

Grey tin ≠ White tin
(Cubic) (Tetragonal)

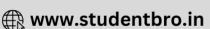
The change of white tin to grey tin is accompanied by increase in volume. This is called tin disease or tin plague.

- 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 $3Na_2O.3Al_2.6SiO_2.2Na_2S$.

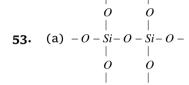




- **45.** (d) In carbon family stability +2 oxidation state increases on moving down the group in the periodic table with an increase in atomic number due to screening effect.
- **46.** (c) Tin is oxidised to meta stannic acid when it is treated with nitric acid.

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

- **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



- **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) $Pb \Rightarrow 11.34 \ g/ml$ 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



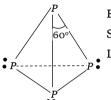
6. (a) White phosphorus is soluble in CS_2 whereas red phosphorus is insoluble in it.

7. (d) $H_4P_2O_7$ pyrophosphoric acid

$$HO - \begin{matrix} O & O \\ \parallel & \parallel \\ P - O - \begin{matrix} P \\ P \end{matrix} - OH \\ \parallel & OH \end{matrix}$$

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_5 is not known because of absence of *d*-orbitals in nitrogen.
- 11. (a,d) P_4 molecule



Bond angle = 60°

Six P-P = Single bonds

Lone pairs = 4

- 12. (b) $NH_4NO_3 \xrightarrow{\Delta} 2H_2O \uparrow + N_2O \uparrow$ Nitrous oxide of auching case
- 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) $NH_4NO_3 \xrightarrow{\Delta} 2H_2O \uparrow + N_2O \uparrow$ $NaNO_3 \xrightarrow{\Delta} NaNO_2 + O_2 \uparrow$ $2AgNO_3(s) \rightarrow 2Ag(s) + 2NO_2(g) + O_2(g)$ 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) N P As Sb BiNon-metals Metalloids Meta
- - 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)
$$2HNO_3 \rightarrow N_2O_5 + H_2O$$

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

 $P_4O_{10} + 10C \xrightarrow{1770K} P_4 + 10CO$

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)
$$(NH_4)_2 Cr_2 O_7 \xrightarrow{\Delta} Cr_2 O_3 + N_2 + 4H_2 O_3$$

31. (d)
$$4HNO_3 \rightarrow 4NO_2 + O_2 + 2H_2O$$

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
$$NH_3 + H_2O \rightarrow NH_4^+ + OH^- \implies NH_4OH$$

(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
$$N_2 + 3H_2 \underbrace{\frac{Fe - Mo}{650 - 800 \, K}} 2NH_3$$

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

$$\begin{matrix} O & OH & O & OH \\ O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} & O & \end{matrix} & O & \end{matrix} & O & \begin{matrix} | \end{matrix} & O & \end{matrix} &$$

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

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

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

46. (d)
$$4NH_3 + 5O_2 \xrightarrow{Pt} 4NO + 6H_2O_3$$

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
 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 $2HNO_3 \rightarrow N_2O_5 + H_2O$

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

57. (b)
$$P_4 O_8$$

$$4x + (-2 \times 8) = 0$$

$$4x - 16 = 0$$

$$x = \frac{16}{4} = +4$$

- (c) NH_2OH 58. x + 2 + (-2) + 1 = 0x + 2 - 2 + 1 = 0x = -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.
- (d) Due to absence of d-orbitals in N atom, it 62. cannot accept electrons from H_2O for hydrolysis of NF_3 .
- (b) NH_3 is most thermally stable hydride. Hence, 63. electrolysis temperature is maximum.
- (a) Phosphorus is kept in water due to it burt at 64. 30°C.
- (c) $BiCl_3 + H_2O \rightarrow BiOCl + 2HCl$ 66.
- (c) When the black ppt. of Bi_2S_3 is dissolved in 67. 50% HNO_3 and a solution of NH_4OH is added. A white ppt. of $Bi(OH)_3$ is obtained.
- 69. (a) Atmospheric nitrogen is inert and unreactive because of very high bond energy (945 kJ/mole).
- (b) Bismuth does not show allotropy other 70. elements show allotropy. Nitrogen $\rightarrow \alpha$ -nitrogen and β -nitrogen (solid crystalline forms) Phosphorus → White, Red and Black forms Arsenic → Yellow and Grey forms Antimony → Yellow and Grey forms
- (a) Nitrogen does not form complexes because of 71. the absence of d-orbitals.
- 72. (a) NH_3 is a strongest base because Lone pair is easily available for donation.
- (b) Hydride AsH_3 SbH_3 74. NH_3 BiH 3

Boling point 238.5 185.5 210.6 254.6 290

- (a) NCl₃ is highly reactive and unstable. Hence it 75. is explosive.
- 76. (b) N_2O_3 As_2O_3 Sb_2O_3 Bi_2O_3 Acidic Oxides Amphoteri Basic

Acidic character decreases down the group

- (c) SbCl 2 is not exists because Vth group elements normally show +3 and +5 oxidation state.
- (b) $NH_4Cl + NaNO_2 \rightarrow NH_4NO_2 + NaCl$ **78.** $NH_4NO_2 \xrightarrow{\text{heat}} N_2 + 2H_2O$

- (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
- (d) $N \equiv N$ bond energy is very high $945 \, kJ \, mol^{-1}$.
- (d) $N_7 \rightarrow 1s^2, 2s^2, 2p^3$ 83. d-orbitals are absent in nitrogen.
- (d) $NH_4NO_3 \xrightarrow{\text{heat}} N_2O + 2H_2O$
- (d) $NH_2OH + HNO_2 \rightarrow H_2 N_2 O_2 + H_2O$
- (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) $2Pb(NO_3)_2 \rightarrow 2PbO + 4NO_2 + O_2$
- (a) $2Pb(NO_3)_2 \rightarrow 2PbO + 4NO_2 + O_2$
- (b) $3Cu + 8HNO_3 \rightarrow 3Cu(NO_3)_2 + 4H_2O + 2NO_3$
- 93. (d) In upper atmosphere NO is formed by lightning flash.

$$N_2 + O_2 \rightleftharpoons 2NO$$

- **96.** (c) $2NO + O_2 \rightarrow 2NO_2$
- (b) $2AgNO_3 \rightarrow 2AgNO_2 + O_2$
- **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

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

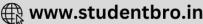
 $HNO_2 + [O] \rightarrow HNO_3$

- **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.
- $N_3H \Rightarrow N_3^- + H^+$ **107.** (d) Hy drazoic acid
- **108.** (a) *d*-orbitals are absent in nitrogen.
- Phosphide ion Chloride ion **109.** (d)

 (P^{3-}) (Cl^{-}) 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. hy pophosph ite
- 117. (c) Both oxidation and reduction (Disproportionation)

Oxidation



B.pt in (K) 238.5 185.5 210.6 254.6

149. (a)
$$H_3PO_2$$

 $3+x-4=0$

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) $Sn + \text{conc. } 4NHO_3 \rightarrow H_2SnO_3 + 4NO_2 + H_2O_3$ Meta stannic acid

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

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

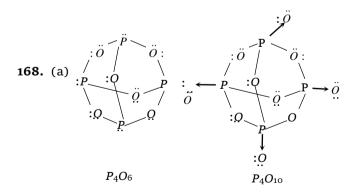
164. (b) Ammomium nitrate is neutral fertilizer.

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. $6Li + N_2 \xrightarrow{450^{\circ} C} 2Li_3N$ (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) $N_2{\cal O}$ on account of stimulating effect on nervous system.

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

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

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

$$H \stackrel{O}{\stackrel{\parallel}{\nearrow}} OH$$
 Only one $-OH$ group

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

127. (b)
$$H_3PO_3$$
 O

$$H \stackrel{\parallel}{\stackrel{\parallel}{\sim}} OH$$

$$OH$$

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.

146. (d) $NH_3 PH_3 AsH_3 SbH_3$



174. (d) Sodium metal in liq. NH_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 Weakbase 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

$$H_3 PO_4 \rightarrow 3H^+ + PO_4^{3-}$$

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) $\frac{NH_3 > PH_3 > AsH_3 > SbH_3 > BiH_3}{2}$

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

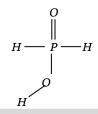
- **204.** (a) Nitrogen forms NH_3 which is most basic.
- **205.** (b) H_3PO_3 is a diabasic acid. It forms two types of salts NaH_2PO_3 and Na_2HPO_3 .
- **206.** (a) $NH_2 CO NH_2 + 2HNO_2 \rightarrow CO_2 + 3H_2O + 2N_2$
- **207.** (c) I II III IV V

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) $Ca_3P_2 + 3H_2O \rightarrow 3Ca(OH)_2 + 2PH_3$
- **213.** (d) $(NH_4)_2Cr_2O_7 \rightarrow N_2 + Cr_2O_3 + 4H_2O_4$
- **214.** (b) B > P > As > Bi As 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_4Cl 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 anhy dride
- **240.** (c) Oxidation number of As in $H_2 As O_4^-$ 2 + x - 8 = -1x - 6 = -1
- **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) $\frac{1}{8}S_8 + 2e^- \rightarrow S^{2-}$

x = 5

- **6.** (c) $2H_2O + 2F_2 \rightarrow 4HF + O_2$
- 9. (b) $O_8 1s^2 2s^2 2p^4$
 - 2 unpaired electrons
- **11.** (a) Element O S Sc Te

- Electronegativity 3.5 2.5 2.4 2.1 2.0
- 13. (b,c) $Na_2SO_3 + H_2SO_4 \rightarrow Na_2SO_4 + H_2O + SO_2$ (S) (aq) (aq) (l) (g)
- **15.** (d) H_2O H_2S H_2Se H_2Te 104.5° 92.1° 91° 90°

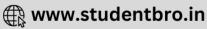
As we go down the group electronegativity decreases due to which repulsion between bonded pairs of electron also decreases. Hence, bond angle decreases.

- **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) $30\frac{\text{silent}}{\text{clectric discharge}} 20_3$
- **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) $K_2Cr_2O_7 + H_2SO_4 + 3SO_2 \rightarrow K_2SO_4 + Cr_2(SO_4)_3 + 3H_2O_4$
- **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) $H_2SO_4 + SO_3 \rightarrow H_2S_2O_7$ (oleum)
- **45.** (b) $2Ag + 2H_2SO_4 \rightarrow Ag_2SO_4 + 2H_2O + SO_2$ Reducing agent Oxidising agent
- **46.** (a) Only dehydrating agent $HCOOH \xrightarrow{H_2SO_4} CO + H_2OOH$





- **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 Fuming H_2SO_4)
- **53.** (d) $S_2 O_7^{2-}$

$$\begin{matrix} O & O \\ \parallel & \parallel \\ -O - S - O - S - O^- \\ \parallel & \parallel \\ O & O \end{matrix}$$

- **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) $HO SO_2 OH + PCl_5 \rightarrow Cl SO_2 OH + POCl_3$

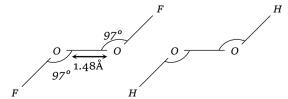
+HCl

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

$$Cl - SO_2 - Cl + 2POCl_3 + 2HCl$$
 Sulphury l 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) 16th group called chalcogens (oxygen family) while *Na* is a Ist 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

$$H_2O + SO_2 \rightarrow H_2SO_3$$

Sulphurus acid

- **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.

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

- **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) $CHCl_3 + \frac{1}{2}O_2 \rightarrow {COCl_2 + HCl}_{\substack{\text{Phosgene or carbonyl} \\ \text{chloride}}} + HCl$ 2.
- 3. (d) Iodine has the least affinity for water and is 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.

- 6. (a) HI is the strongest reducing agent among 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.
- (a) HF is the weakest acid. Since it is unable to 17. give H^+ ions which are trapped in H-Bonding.
- (b) Hydride -HCl HBrHI 18. HF189 206 B.pt (in K) - 293 238 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.
- (a) +5 22. +3 $HClO_4 > HClO_3 > HClO_2 > HClO$

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

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

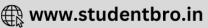
$$2KHSO_4 + MnSO_4 + 2H_2O + Br_2$$

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

Boiling -34.659.5 185.2 $(^{\circ}C)$ 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
- (d) As number the atomic 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. $H_2SO_4 + 2HI \rightarrow 2H_2O + I_2 + SO_2 \uparrow$
Violet

74. (b)
$$2NaI + Cl_2 \rightarrow 2KCl + I_2$$

 $CCl_4 + I_2 \rightarrow Violet colour$

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 $(SCN)_2$ Thiocyanogen

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

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

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

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

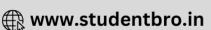
Cl > S > P.

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

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) $2NaOH + Cl_2 \rightarrow NaCl + NaOCl + H_2O$ $\downarrow \qquad \qquad \downarrow \qquad \downarrow$ $Na^+ + Cl^- Na^+ + OCl^-$
- 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
- **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 Cl_2 is less electronegative then 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) $3KClO_3 + 3H_2SO_4 -$

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

- **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 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^- ClO₂ 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 the (Cl_2O_7) is anhydride of perchloric acid. $2HClO_4 \xrightarrow{\Delta} Cl_2O_7 + H_2O$
- **187.** (c) I_2 forms complex ion I_3^- in KI solution due to which it dissolves in it.

Noble gases

- (a) Gas -3. H_2 O_2 He N_2 Mol. mass - 2 32 4
- (c) Helium (In Greek Helios = Sun) 5.
- 6. the noble gases are colourless and odourless gases. 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.
- (c) Except He, all other noble gases are adsorbed 7. by coconut charcoal at low temperatures. The extent of adsorption increases as the atomic size of the noble gas increases.

- (b) An oxygen-helium mixture is used artificial 10. 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".
- (c) XeF_2 , $XeOF_2$. XeF_4 , $XeOF_4$. XeF_6 , XeO_3 .
- (c) Gas (Abundance in air by volume (ppm))

Helium	5.2
Neon	18.2
Argon	93.4
Krypton	1.1
Xenon	0.09

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

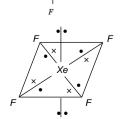
$$F$$
 Xe
 F

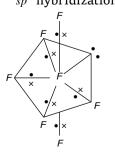
- (b) Partial hydrolysis; $XeF_4 + H_2O \rightarrow XeOF_2 + 2HF$ Complete hydrolysis; $2XeF_4 + 3H_2O \rightarrow Xe + XeO_3 + F_2 + 6HF$
- (d) He is least polarizable because of small atomic size
- 27. (a) Rn because it is radioactive element obtained by the disintegration of radium $_{88}$ Ra $^{206} \rightarrow _{86}$ Rn $^{202} +_{2}$ He 4
- **30.** (c) $1s^2 2s^2 2p^6 \rightarrow \text{Neon}$

(d)

37.

- (a) $XeF_6 + 3H_2O \rightarrow XeO_3 + 6HF$ 35.
- (c) Xe > Kr > Ar > Ne > HeSolubility in decreasing order.
 - sp³ hybridization



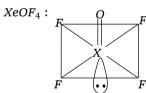


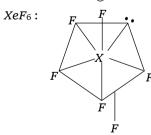
 sp^3d^2 hybridization sp^3d^3 hybridization

- (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. $Xe + F_2 \xrightarrow{Ni \text{ tube}} XeF_2$; $Xe + 2F_2 \xrightarrow{673 \text{ K}} XeF_4$

 $Xe + 3F_2 \xrightarrow{523-573 K} XeF_6$ 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) XeO₃:





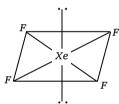
- (d) Neil Bartlett prepared first noble gas 51. compound. Xenon hexafluoroplatinate (IV).
- (d) He, Ne, and Kr all are found in very little 53. amount in atmosphere, so all are called rare
- (c) Helium is twice as heavy as hydrogen, its 54. lifting power is 92% of that of hydrogen. Helium has the lowest 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. $XeOF_{2}$

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

- **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. metaaluminate (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) $ZnO + BaO \xrightarrow{1100^{\circ}C} BaZnO_{?}$
- **30.** (b) There are no free electron in N_2O_4 , so it is dimagnetic in nature.
- 31. (b) $CaSO_4.2H_2O \xrightarrow{120^{\circ}C} CaSO_4.\frac{1}{2}H_2O + 1\frac{1}{2}H_2O$ 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) $NH_4NO_2 \rightarrow N_2 + 2H_2O$ $2NaN_3 \rightarrow 2Na + 3N_2$ $(NH_4)_2 Cr_2 O_7 \rightarrow N_2 + Cr_2 O_3 + 4H_2 O$
- **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. $I_2O_7 \& I_2O_5$ 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 $Fe_2O_3 + Al$

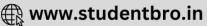
- **68.** (d) The colour of liquid O_2 is pale blue.
- **69.** (a) Helium is not soluble in blood even under high pressure, a mixture of 80% helium and 20% oxygen is used instead of ordinary air by sea divers for respiration.
- **70.** (a) $Mg_3N_2 + 6H_2O \rightarrow 3Mg(OH)_2 + 2NH_3$
- **71.** (a)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

Assertion & Reason

- (b) Sulphate is estimated as BaSO₄ 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.
- (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 $NaAlO_2$.







 $Al(OH)_3 + NaOH \rightarrow \underbrace{NaAlO_2}_{\text{Sodium meta aluminate}} + 2H_2O$

- **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.

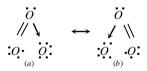
- 19. (d) Anhydrous BaO_2 is not used for preparing H_2O_2 because it reacts with H_2SO_4 and the reaction ceases after some time due to formation of $BaSO_4$ on BaO_2 . Therefore, assertion and reason both are false.
- **20.** (d) Inorganic benzene, borazine is highly reactive while benzene is much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

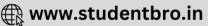
- **27.** (c) Assertion is true but reason is false. Due to high polarizing power of $Li^+, LiCl$ is a covalent compound.
- **28.** (c) Assertion is true but reason is false. *Be* has fully filled $2s^2$ orbital which gives a relatively more stable electronic configuration.
- **29.** (a) Both assertion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



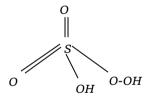


- **40.** (b) Both assertion and reason are true but reason is not the correct explanation of assertion. In case of NI_3 , the lone pair moment adds on the resultant of the N-I moments but in case of NF_3 , the lone pair moment on partly cancels the resultant N-F moments.
- 41. (a) Both assertion and 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^3 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*₃ 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 waxlined 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 space lattice.
- 59. (d) Both assertion and reason are false.





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.

