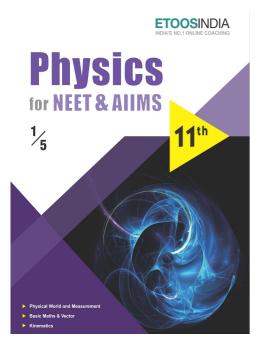
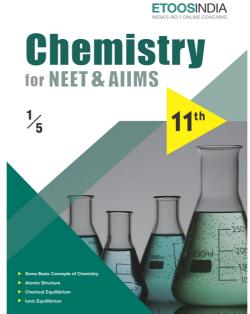
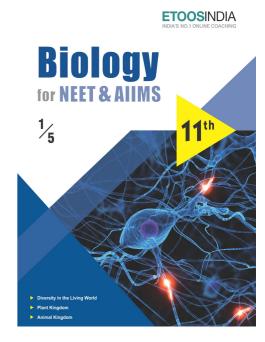
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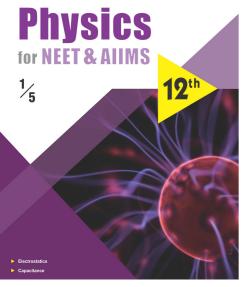
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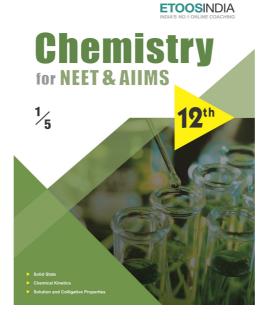


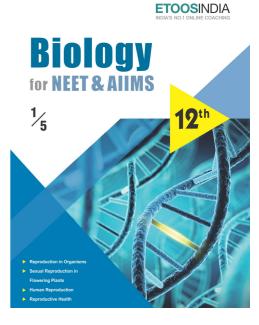












ETOOS Comprehensive Study Material For NEET & AIIMS

CHAPTER 0

### **P-BLOCK ELEMENTS**

if all the elements are arranged in the order of atomic weights, a periodic repitition is obtained. This is expressed by the law of periodicity.

"DMITRI MENDELEEV"

#### **INTRODUCTION**

he elements is which last electron enters into p-subshell are called as p-block elements. The number of p-orbitals are three and, therefore, the maximum number of electrons that can be accommodated in a set of p-orbitals is six, hence p-block contains six groups. The general electronic configuration of these elements is ns<sup>2</sup>np<sup>1-6</sup> (except for He).

p-block elements can be either metals, metalloids or non-metals. Maximum oxidation state shown by p-block elements is equal to the sum of valence electrons (i.e., sum of s-and p-electrons) or the group number minus 10 known as group oxidation state.

Absence of d-orbitals in the elements of second period and presence of d- and f-orbitals in rest of elements have significant effects on the properties as a result lower oxidation state will be more stable. This effect is known as **inert pair effect**.

- (c) **Hybridisation**: If a lone pair participates in back bonding then it is not considered in hybridisation. Ex.: B<sub>3</sub>N<sub>3</sub>H<sub>6</sub> (inorganic benzene or borazene or borazele)
- Hybridisation of B as well as  $N = sp^2$

- Inorganic benzene is more reactive than organic benzene as in it the bonds are polar, although over all molecule is non polar.
- (d) If back bonding is present then tendency to form dimer or polymer decreases.  $Ex.: BF_3, BeF_2$
- (2)  $p\pi$ -d $\pi$  back bonding:

it is used to explain following observations:

(a) Hybridisation

Ex. Trimethyl amine (CH<sub>3</sub>)<sub>3</sub>N

- sp<sup>3</sup> hybrid (N)
- trigonal pyramidal
- Lewis base (due to presence of lp)
- (b) Acidic strength

- · No back bonding
- Less acidic

#### Trisilyl amine

$$(SiH_3)_3 N$$

- sp<sup>2</sup> hybrid (N)
- trigonal planar
  - Not Lewis base
- Bond angle increases

$$\begin{array}{c} SiH_{\scriptscriptstyle 3} - \overset{.}{O} \stackrel{!}{=} H \\ \text{Silyl alcohol} \end{array}$$

- Back bonding present in conjugate base
- More acidic

#### (II) Dimerisation / Polymerisation

# Types of Dimerisation 3C—2e $B_2H_6$ $(BeCl_2)_2$ $(BeH_2)_0$ $(BeH_2)_n$ $Al_2(CH_3)_6$ $Ga_2(CH_3)_6$ $(ICl_3)_2$



#### **ETOOS KEY POINTS**

Reactivity towards halogen:

ement in pure state	Type of bonding	<b>Meling Point</b>
C	Covalenet	4100°C
Si	Covalent	1420°C
Ge	Covalnet	945°C
Sn	Metallic	232°C
Ph	Metallic	327°C

On moving down the group from carbon to lead stability of +4 oxidation state decreases while stability of +2 oxidation state increases andd hence decreases oxidising power decreases down the group due to inert pair effect.

These elements form two types of hallides –  $MX_2$  and  $MX_4$ . Most of the  $MX_4$  are covalent.  $SnF_4$  and  $PbF_4$  are ionic in nature.

Thermal stability decreases with increasing atomic size of molecular mass of tetrahalide or due to decreasing polarity.

$$CX_4 > SiX_4 > GeX_4 > SnX_4 > PbX_4$$
 and  $CF_4 > CCl_4 > CBr_4 > Cl_4$ 

In these compounds:

Hybridisation  $\rightarrow sp^3$ 

Geometry → Regular tetrahedral

Polarity  $\rightarrow$  NON-POLAR

Bond angle  $\rightarrow$  109° 28′

 $PbI_4$  is not stable as  $I^-$  is strong reducing agent which reduces  $Pb^{+4}$  to  $Pb^{+2}$  and also stability of +4 oxidation state of Pb is lesser than +2 state.

Sodium Zeolite [Na<sub>2</sub>Al<sub>2</sub>Si<sub>2</sub>O<sub>8</sub>] / [Na<sub>2</sub>O.Al<sub>2</sub>O<sub>3</sub>.2SiO<sub>2</sub>]

- (i) It is a 3-D silicate
- (ii) It is used in
  - (a) For softening of hard water
  - (b) For cracking of hydrocarbon & isomerisation

Ex. ZSM-5 (Zeolite) is used to convert ethyl alcohol into petrol.

#### Etoos Tips & Formulas

#### **BORON FAMILY**

(1) Amorphous boron of low purity (called moissan boron) is obtained by reducing B<sub>2</sub>O<sub>3</sub> with Mg or Na at a high temperature. It is 95–98% pure (being contaminated with metal borides), and is black in colour.

$$Na_2[B_4O_5(OH)_4].8H_2O$$
 acid  $H_3BO_3$  heat  $B_2O_3$  Mg or  $Na$   $2B+3$  MgO

(2) BORANES

Boranes are boron hydrogen compounds with general molecular formula  $B_nH_{n+4}$  or  $B_nH_{n+6}$ . They are electron deficient compounds.

(3) ALUM

Alums are double sulphates with their general formula  $R_2SO_4.M_2(SO_4)_3.24H_2O$  where  $R = monovalent \ radical \ like \ Na^+, K^+, NH_4^+ \ and M = Trivalent \ radical \ like \ Al^{+3}, Cr^{+3}. Fe^{+3}.$ 

#### **CARBON FAMILY**

- (1) Carbon is found in nature in various allotropic forms which are:
  - (i) Crystalline Form: Diamond, Graphite, Fullerenes
  - (ii) Amorphous Form: Coal
- (2) SILICON (Si)

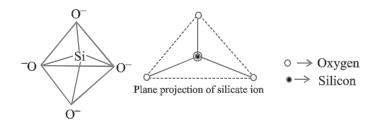
Silicon is the second most abundant (27.2%) element after oxygen (45.5%) in the earth's crust. It does not occur free in nature but in the combined state, it occurs widely in form of silica and silicates. All mineral rocks, clays and soils are built of silicates of magnesium, aluminium, potassium or iron. Aluminium silicate is however the most common constituent of rocks and clays.

Silica is found in the free state in sand, flint and quartz and in the combined state as silicates like

- (a) Feldspar  $K_2O.Al_2O_3.6SiO_2$
- (b) Kaolinite Al<sub>2</sub>O<sub>3</sub>. 2 SiO<sub>2</sub>. 2H<sub>2</sub>O
- (c) Asbestos CaO. 3MgO.4SiO<sub>2</sub>

**Silicates** are also important:

Silicates have basic unit of SiO<sub>4</sub><sup>4-</sup>, each silicon atom is bonded with four oxide ions tetrahedrally.



#### **SOLVED EXAMPLE**

- Ex. 1 By adding gypsum to cement
  - (A) Setting time of cement becomes less
  - (B) Setting time of cement increases
  - (C) Colour of cement becomes light
  - (D) Shining surface is obtained
- Ans. (B) Raw materials for cement-limestone, clay, gypsum, Cement is a dirty greyish heavy powder containing calcium aluminates and silicated.

Gypsum (CaSO<sub>4</sub>.5H<sub>2</sub>O) is added to the components to increases the setting time of cement to that it gets sufficiently hardened. Setting of cement is an exothermic process and involves hydration of calcium aluminated and silicates.

- Ex. 2 Silcon has a strong tendency to form polymers like silicones. The chain length of silicon polymer can be controlled by adding
  - (A) MeSiCl,
- (B) Me,SiCl,
- (C) Me,SiCl
- (D) Me<sub>4</sub>Si
- Ans. (C) Silicon has a strong tendency to form polymers like silicones. The chain length of silicon polymer can be controlled by adding Me<sub>3</sub>SiCl which block the ends as shown below

- Ex. 3 Quartz is extensively used as a piezoelectric material. it contains.....
  - (A) Pb
- (B) Si

(C) Ti

- (D) Sn
- Ans. (B) Quartz, cristobalite and tridymite are some of the crystalline forms of silica and they are intercovertable at suitable temperature. Quartz is edxtensively used as a piezoelectric material.
- Ex. 4 The most commonly used reducing agent is
  - (A) AlCl<sub>3</sub>
- (B) PbCl,
- (C) SnCl,
- (D) SnCl<sub>2</sub>
- Ans. (D) Reducing agents are those substance which resduced other subastance and it self oxidises

  In SnCl<sub>2</sub>, Sn exists in +2 oxidation state, thus, acts

$$SnCl_2 + 2FeCl_3 \longrightarrow 2FeCl_2 + SnCl_4$$

as a strong reducing agent i.e.,

$$SnCl_2 + 2CuCl_2 \longrightarrow 2CuCl + SnCl_4$$

- Ex. 5 Which of the following piars of ions are isolectonic and isostructural
  - (A)  $CO_{2}^{2-}$ ,  $NO_{2}^{-}$
- (B)  $ClO_3^-, CO_3^{2-}$
- $(\mathbb{C}) SO_3^{2-}, NO_3^{-}$
- (**D**)  $ClO_3^-, SO_3^{2-}$
- Ans. (A) Compounds having same value of total number of electrons are known as isolectronic.

For CO<sub>2</sub><sup>2-</sup>

For NO<sub>2</sub>

Total number of electons

Total number of electrons

$$=6+8\times3+2$$

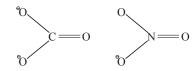
$$= 7 + 8 \times 3 + 1$$

$$=6+24+2$$

$$=7+25$$

$$=32$$

$$=32$$



Hence,  $CO_3^{2-}$  and  $NO_3^{-}$  are isolectronic. These two ions have similar structure so they are isostructural. Both have triangular planar structure as in both the species carbon and nitrogen are  $sp^2$  hybridised.

#### SINGLE OBJECTIVE NEET LEVEL Exercise # 1 The most acidic of the following compounds is 1. 10. Which of the following is most acidic $(A) P_2O_3$ (B) Sb<sub>2</sub>O<sub>3</sub> (A) Na<sub>2</sub>O (B) MgO $(C) B_2O_2$ $(D) As_2O_2$ (C) Al,O, (D) CaO 2. Identify the statement that is not correct as far as 11. When orthoboric acid (H<sub>2</sub>BO<sub>2</sub>) is heated, the resistructure of diborane is concerned due left is (A) There are two bridging hydrogen atoms in (A) Metaboric acid (B) Boron diborane (D) Borax (C) Boric anhydride (B) Each boron atom forms four bonds in diborane (C) The hydrogen atoms are not in the same plane 12. Silicon dioxide is formed by the reaction of in diborane (A) SiCl<sub>4</sub> + 2H<sub>2</sub>O (B) SiO<sub>2</sub> + 4HF (D) All B - H bonds in diborane are similar (C) SiO, + NaOH (D) SiCl, + NaOH Soft heavy metal melts at and is used in making 3. 13. Which alkali metal carbonate decomposes on heatheat sensitive thermometers the metal is ing to liberate CO, gas (A) Galium (B) Sodium (B) CaCO, (A) Li,CO, (C) Potassium (D) Caesium (D) Al<sub>2</sub>CO<sub>2</sub> (C) Na<sub>2</sub>CO<sub>3</sub> 4. Which of the following is formed when aluminium oxide and carbon is strongly heated in dry chlorine 14. Which of the following gives propyne on hydrolysis (A) Aluminium chloride $(A) Al_{4}C_{2}$ $(B) Mg_2C_2 \qquad (C) B_4C$ (B) Hydrate aluminium chloride (C) Anhydrous aluminium chloride 15. Which one of the following statements is not (D) None of these correct 5. Which metal burn in air at high temperature with (A) Zinc dissolves in sodium hydroxide solution the evolution of much heat (B) Carbon monoxide reduces iron (III) oxide to iron (A) Cu (B) Hg (C) Mercury (II) iodide dissolves in excess of (C) Pb (D) Al potassium iodide solution (D) Tin (IV) chloride is made by dissolving tin so-Aluminium hydroxide is soluble in excess of sodium 6. hydroxide forming the ion lution in concentrated hydrochloric acid $(A) AlO_{2}^{+3}$ (B) $AlO_2^{-3}$ 16. In laboratory silicon can be prepared by the reaction (C) AlO, (D) AlO<sub>2</sub> (A) By heating carbon in electric furnace 7. Boron form covalent compound due to (B) By heating potassium with potassium (A) Higher ionization energy dichromate (B) Lower ionization energy (C) Silica with magnesium (C) Small size (D) None of these (D) Both (A) and (C) 17. Which of the following is the correct statement for 8. In diborane, the two H–B–H angles are nearly red lead **(B)** $95^{\circ}$ , $120^{\circ}$ (A) $60^{\circ}$ , $120^{\circ}$ (A) It is an active form of lead (C)95°, 150° (D) 120°, 180° (B) Its molecular formula is Pb<sub>2</sub>O<sub>3</sub> 9. Which of the following is a non-metal (C) It decomposes into Pb andm CO, (A) Gallium (B) Indium

(C) Boron

(D) Aluminium

(D) It decomposes into PbO and O,

## Exercise # 2

#### SINGLE OBJECTIVE

#### AIIMS LEVEL

1.	Boric acid polymerizes d (A) The presence of hyd (B) Its acidic nature		11.	Silicones have the general (A) SiO <sub>4</sub> <sup>4-</sup> (C) (R <sub>2</sub> SiO) <sub>n</sub>	formula - (B) Si <sub>2</sub> O <sub>7</sub> <sup>6-</sup> (D) (SiO <sub>3</sub> ) <sub>n</sub> <sup>2-</sup>
	<ul><li>(C) Its geometry</li><li>(D) Its monobasic nautre</li></ul>		12.	In which of the following thing –	-
2.	Aluminium is obtained b (A) Reduction of Al <sub>2</sub> O <sub>3</sub> w			<ul><li>(A) Diamond</li><li>(C) Dimethylamine</li></ul>	<ul><li>(B) Graphite</li><li>(D) Trisilylamines</li></ul>
	(B) Electrolysis of Al <sub>2</sub> O <sub>3</sub> (C) Reduction of Al <sub>2</sub> O <sub>3</sub> w (D) Heating cryolite and a	dissolved in Na <sub>3</sub> AlF <sub>6</sub> ith chromium	13.	Glass or silica soluble in – (A) HClO <sub>4</sub> (C) Aqua-regia	(B) HF (D) H <sub>2</sub> SO <sub>4</sub>
3.	In thermite welding, alun (A) A solder (C) An oxidising agent	ninium acts as –  (B) A flux  (D) A reducing agent	14.	The species present in solution when CO <sub>2</sub> dissolved in water are –  (A) CO <sub>2</sub> ,H <sub>2</sub> CO <sub>3</sub> , HCO <sub>3</sub> <sup>-</sup> ,CO <sub>3</sub> <sup>2-</sup> (B) H <sub>2</sub> CO <sub>3</sub> ,CO <sub>3</sub> <sup>2-</sup> (C) CO <sub>3</sub> <sup>2-</sup> ,HCO <sub>3</sub> <sup>-</sup> (D) CO <sub>2</sub> ,H <sub>2</sub> CO <sub>3</sub>	
4.	to red heat is –	d when boric acid is heated			
5.	(A) Metaboric acid (C) Boron oxide  Which of the following ca	(B) Tetraboric acid (D) Pyroboric acid an be detected by the borax-	15.	P <sub>2</sub> O <sub>5</sub> is used extensively a (A) Dehydrating agent (C) Reducing agent	s a –  (B) Catalytic agent  (D) Preservative
	bead test ? (A) Ni <sup>2+</sup> (C) Pb <sup>+2</sup>	(B) Co <sup>2+</sup> (D) Both (A) & (B)	16.	The number of molecules vert one molecules of P <sub>2</sub> acid is –	s of water needed to con-
6.	The hydrides of boron ar (A) Boron hydrogen con	re called		(A) 2 (C) 4	(B) 3 (D) 5
	<ul><li>(B) Hydrogen borides</li><li>(C) Boranes</li><li>(D) Hydroboric acids</li></ul>		17.	Producer gas is a mixture (A) CO and $N_2$ (C) CO and $H_2$	of – $ (B) CO2 and H2 $ $ (D) CO2 and N2 $
7.	an alum ? (A) K <sub>2</sub> SO <sub>4</sub> .Al <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub> .24H		18.	Which variety of glass is optical glasses? (A) Sodium glass	(B) Flint glass
	(B) K <sub>2</sub> SO <sub>4</sub> .Cr <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub> .24H <sub>2</sub> (C) Na <sub>2</sub> SO <sub>4</sub> .Fe(SO <sub>4</sub> ) <sub>3</sub> .24H (D) CuSO <sub>4</sub> .Al <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub> .24H	_O	19.	(C) Ground glass  The colour imparted by C is –	(D) Quartz o(II) compounds to glass
8.	Higher percentage of car (A) Anthracite			(A) Green (C) Yellow	(B) Deep–Blue (D) Red
9.	(C) Bituminous  From B <sub>2</sub> H <sub>4</sub> , all the following	(D) Peat  ng can be prepared except –	20.	In warfare smoke screens (A) PH <sub>3</sub>	(B) CaC <sub>2</sub>
	$(A) B_2 O_3$ $(C) B_2 (CH_3)_6$	(B) H <sub>3</sub> BO <sub>3</sub> (D) NaBH <sub>4</sub>	21.	(C) P <sub>2</sub> O <sub>5</sub> In Haber's process for the r	(D) COCl <sub>2</sub> manufacture of ammonia,
10.	The product formed in the $BCl_3 + H_2O \longrightarrow Product$ (A) $H_3BO_3 + HCl$ (C) $B_2H_6 + HCl$			the catalyst used is –  (A) Finely divided nickel  (B) Finely divided molybo  (C) Finely divided iron  (D) Finely divided platinus	

(D) Finely divided platinum

#### Exercise #3

#### PART - 1

#### MATRIX MATCH COLUMN

1. Match the reactions listed in column-I with characteristic(s) / type of reactions listed in column-II.

Column-I

Column-II

(A)  $BBr_3 + H_2 \longrightarrow B$ 

- (p) Borax bead test
- (B)  $Na_2B_4O_7$ .  $10 H_2O + CuSO_4 \rightarrow Cu(BO_2)_2$
- (q) Reduction

 $(\mathbb{C}) \text{ AlCl}_3 + \text{H}_2\text{O} \longrightarrow \text{HCl}$ 

(r) White fumes

(D)  $Cr_2O_3 + Al \longrightarrow Cr$ 

- (s) Hydrolysis
- 2. Match the reactions listed in column-I with characteristic(s) / type of reactions listed in column-II.

Column-I

Column-II

 $(A) Al_2(C_2)_3 + H_2O \longrightarrow$ 

(p) One of the products contains both  $\sigma$  and  $\pi$  bonds

(B)  $CH_2(COOH)_2 + P_4O_{10} \longrightarrow$ 

(q) Hydrolysis

(C)  $CH_3 SiCl_3 + H_2O \longrightarrow$ 

(r) Dehydration

(D)  $SnCl_2.2H_2O \xrightarrow{on standing}$ 

- (s) complex crosslinked polymer
- 3. Match the reactions listed in column-I with the product(s) listed in column-II.

Column-I

Column-II

 $(A) B_2O_3 + H_2O$ 

(p) H<sub>3</sub>BO<sub>3</sub>

 $(B) B_2 H_6 + H_2 O$ 

(q) H,

 $(C) B_3 N_3 H_6 + H_2 O$ 

(r) HCl

(7) 3 3 6 2

(s) NH,

(D)  $BCl_3 + H_2O$ 

- (t) N,
- 4. Match the type of silicates listed in column-I with characteristic(s) listed in column-II.

Column-I

Column-II

(A) Cyclic silicates

(p) Tetrahedral hybridisation.

(B) Single chain silicates

(q) Si–O bonds are 50% ionic and 50% covalent.

(C) Pyro silicates

- (r) General formula is  $(SiO_3)_n^{2n-}$
- (D) Sheet silicates (two dimensional)
- (s) Two oxygen atoms per tetrahedron are shared.
- 5. Match the materials listed in column-I with type of silicates listed in column-II.

Column-I

Column-II

(A) Spondumene

(p) Two dimensional sheet silicates

(B) Thorteveitite

(q) Pyrosilicates

(C) Kaolin

(r) Chain silicates

(D) Quartz

(s) Three dimensional sheet silicates.

#### Exercise # 4

#### PART - 1

#### PREVIOUS YEAR (NEET/AIPMT)

1. Which one of the following arrangements does not truly represent the property indicated against it?

[CBSE AIPMT 2000]

- (A)  $Br_2 < Cl_2 < F_2$ , Oxidising power
- (B)  $Br_2 < Cl_2 < F_2$  Electronegativity
- (C)  $Br_2 < F_2 < Cl_2$  Electron affinity
- (D)  $Br_2 < Cl_2 < F_2$  Bond energy
- 2. Among the following the electron deficient compound is [CBSE AIPMT 2000]
  - (A) BCl,

(B) CCl<sub>4</sub>

(C) PCl<sub>5</sub>

- (D) BeCl,
- Nitogen forms  $N_2$ , but phosphorus when form  $P_2$  readily converted into  $P_4$  reason is

[CBSE AIPMT 2001]

- (A) triple bond present between phosphorus atom
- (B)  $p\pi$  - $p\pi$  bonding is weak
- (C)  $p\pi$  bonding is strong
- (D) multiple bond form easily
- 4. Which reaction is not feasible ?[CBSE AIPMT 2002]
  - (A)  $2Kl + Br_2 \longrightarrow 3KBr + l_2$
  - (B)  $2KBr + l_2 \longrightarrow 2KCl + Br_2$
  - (C)  $2KBr + Cl_2 \longrightarrow 2KCl + Br_2$
  - (D)  $2H_2O + 2F_2 \longrightarrow 4HF + O_2$
- 5. Which of the following statements is true?

[CBSE AIPMT 2002]

- (A) Silicon exhibits 4 coordination number in its compounds
- (B) Bond energy of F, is less than Cl,
- (C) Mn(III) oxidation state is more stable than Mn(II) in aqueous state
- (D) Elements of 15<sup>th</sup> group shows only +3 and +5 oxidation states
- 6. In borax bead test which compound is formed?

[CBSE AIPMT 2002]

- (A) Ortho borate
- (B) Meta borate
- (C) Double oxide
- (D) Tetra Borate
- 7. Zn gives H<sub>2</sub> gas with H<sub>2</sub>SO<sub>4</sub> and HCl but not with HNO<sub>3</sub> because [CBSE AIPMT 2002]
  - (A) Zn act as oxidising agent when react with HNO<sub>3</sub>
  - (B) HNO<sub>3</sub> is weaker acid than H<sub>2</sub>SO<sub>4</sub> and HCl
  - (C) In electrochemical series Zn is placed above hydrogen
  - (D) NO<sub>2</sub> is reduced in preference to hydronium ion

The oxidation states of sulphur in the anious  $SO_3^{2-}$ ,  $S_2O_6^{2-}$  and  $S_2O_6^{2-}$  follows the order

[CBSE AIPMT 2003]

(A) 
$$S_2O_4^{2-} < S_2O_6^{2-} < SO_3^{2-}$$

(B) 
$$S_2O_6^{2-} < S_2O_4^{2-} < SO_3^{2-}$$

(C) 
$$S_2O_4^{2-} < SO_3^{2-} < S_2O_6^{2-}$$

(D) 
$$SO_3^{2-} < S_2O_4^{2-} < S_2O_6^{2-}$$

- Which one of the following compounds is not a protonic acid [CBSE AIPMT 2003]
- (A) HOCl is a stronger acid than HOBr
- (B) HF is a stronger acid than HCl
- (C) Among hailde ions. iodide is the most powerful reducing agent
- (D) Fluorine is the only halogen that does not show a variable oxidation state
- 10. Which one of the following compounds is not a protonic acid? [CBSE AIPMT 2003]
  - $(A) SO(OH)_2$
- (B) SO<sub>2</sub>(OH),
- $(\mathbb{C})$  B(OH),

11.

- (D) PO(OH),
- Which one of the following statements about the zeolites is false? [CBSE AIPMT 2004]
  - (A) They are used as cation exchangers
  - (B) They have open structure which enables them to take up small molecules
  - (C) Zeolites are aluminosilcates having three dimensional network
  - (D) Some of the  $SiO_4^{4-}$  units are replaced by  $AIO_4^{5-}$  and  $AIO_6^{9-}$  ions in zeolites
- 12. Which is the best description of behaviour of bromine in the reaction given below?

 $H_2O + Br_2 \longrightarrow HBr + HoBr$  [CBSE AIPMT 2004]

- (A) Only oxidised
- (B) Only reduced
- (C) Both oxidised and reduced
- (D) Only proton accepted

#### MOCK TEST

#### STRAIGHT OBJECTIVE TYPE

1.	Ozone with KI soluti (A) Cl <sub>2</sub>	on produces (B) I <sub>2</sub>	(C) HI	(D) IO <sub>3</sub>		
2.	The gases respectivel (A) O <sub>3</sub> , CH <sub>4</sub>	y absorbed by alkaline py (B) O <sub>2</sub> , O <sub>3</sub>	rogallol and oil of cinnamor (C) SO <sub>2</sub> , CH <sub>4</sub>	n is (D) N <sub>2</sub> O, O <sub>3</sub>		
3.	Ozone turns trimethyl (A) Green	paper (B) Violet	(C) Red	(D) Black		
4.	Ozone is obtained fro (A) By oxidation at hi (C) By silent electric of	gh temperature	(B) By oxidation usin (D) By conversion at	•		
5.	<ul><li>(A) It is harmful beca</li><li>(B) It is beneficial be</li><li>(C) It is beneficial be</li></ul>	cause ozone cuts out the	living organism can proceed faster in the presultraviolet radiation of the su			
6.	The correct order of the	he thermal stability of hyd	drogen halides (H -X) is			
	(A) $HI > HBr > HCl$	> HF	(B) $HF > HCl > HB$	r > HI		
	( $\mathbb{C}$ ) HCl < HF < HBr	< HI	(D) $HI > HCI < HF$	< HBr		
7.	Phosgene is the comm (A) Carbonyl chloride (C) Phosphorus oxycl	;	<ul><li>(B) Phosphine</li><li>(D) Phosphorus trich</li></ul>	<ul><li>(B) Phosphine</li><li>(D) Phosphorus trichloride</li></ul>		
8.	The solubility of iodin (A) Alcohol	ne in water increases in the (B) Chloroform	ne presence of  (C) Sodium hydroxid	e (D) Potassium iodide		
9.	When thiosulphate io	When thiosulphate ion is oxidised by iodine, which one of the following ion is produced				
	(A) $SO_3^{2-}$	(B) SO <sub>4</sub> <sup>2-</sup>	(C) S <sub>4</sub> O <sub>6</sub> <sup>2-</sup> (Tetrathio	onate) (D) $S_2O_6^{2-}$		
10.	Which one of the foll (A) Xe	owing noble gases is the (B) Ar	least polarizable (C) Ne	(D) He		
11.	Which one of the foll (A) Rn	owing noble gases is not (B) Kr	found in the atmosphere (C) Ne	(D) Ar		
12.		n blood than nitrogen at litrogen ble with oxygen	deep sea divers because high pressure			
13.	Which of the followin (A) Ar is used in election (C) Half life of Rn is		(B) Kr is obtained du	uring radioactive disintegration ducing very low temperature		
14.	Aqueous solution of orthoboric acid can be titrated against sodium hydroxide using phenolphthalein indicate only in the presence of :					
	(A) trans-glycerol	(B) catechol	(C) cis-glycerol	(D) both (B) and (C)		

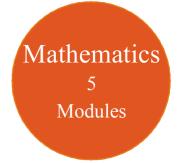
# 11th Class Modules Chapter Details

Physics
5
Modules

1. Oscillations

2. Waves

Chemistry
5
Modules



3. Plant Growth and Development

5. Breathing & Exchange of Gases

1. Body Fluids & Its Circulation

2. Excretory Products & Their

3. Locomotion & Its Movement

4. Neural Control & Coordination5. Chemical Coordination and

**4.** Digestion & Absorption

**Module-5** 

Elimination

Integration

PHYSICS	CHEMISTRY	BIOLOGY
Module-1	Module-1(PC)	Module-1
<ol> <li>Physical World &amp; Measurements</li> <li>Basic Maths &amp; Vector</li> <li>Kinematics</li> </ol>	<ol> <li>Some Basic Conceps of Chemistry</li> <li>Atomic Structure</li> <li>Chemical Equilibrium</li> </ol>	<ol> <li>Diversity in the Living World</li> <li>Plant Kingdom</li> <li>Animal Kingdom</li> </ol>
Module-2  1. Law of Motion & Friction 2. Work, Energy & Power  Module-3	<ul> <li>4. Ionic Equilibrium</li> <li>Module-2(PC)</li> <li>1. Thermodynamics &amp; Thermochemistry</li> <li>2. Redox Reaction</li> <li>3. States Of Matter (Gaseous &amp; Liquid)</li> </ul>	<ul> <li>Module-2</li> <li>1. Morphology in Flowering Plants</li> <li>2. Anatomy of Flowering Plants</li> <li>3. Structural Organization in Animals</li> <li>Module-3</li> </ul>
<ol> <li>Motion of system of particles &amp; Rigid Body</li> <li>Gravitation</li> <li>Module-4</li> <li>Mechanical Properties</li> </ol>	Module-3(IC)  1. Periodic Table 2. Chemical Bonding 3. Hydrogen & Its Compounds 4. S-Block	1. Cell: The Unit of Life 2. Biomolecules 3. Cell Cycle & Cell Division 4. Transport in Plants 5. Mineral Nutrition
of Matter 2. Thermal Properties of Matter  Module-5	Module-4(OC)  1. Nomenclature of Organic Compounds	Module-4  1. Photosynthesis in Higher Plants 2. Respiration in Plants

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2. Isomerism

Module-5(OC)

3. General Organic Chemistry

1. Reaction Mechanism

3. Aromatic Hydrocarbon

4. Environmental Chemistry &

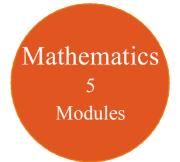
Analysis Of Organic Compounds

**2.** Hydrocarbon

# 12<sup>th</sup> Class Modules Chapter Details

Physics 5 Modules

Chemistry 5 Modules



2. Biodiversity and Conservation

3. Environmental Issues

PHYSICS	CHEMISTRY	BIOLOGY
Module-1	Module-1(PC)	Module-1
<ol> <li>Electrostatics</li> <li>Capacitance</li> <li>Module-2</li> <li>Current Electricity</li> </ol>	<ol> <li>Solid State</li> <li>Chemical Kinetics</li> <li>Solutions and Colligative Properties</li> </ol> Module-2(PC)	<ol> <li>Reproduction in Organisms</li> <li>Sexual Reproduction in Flowering Plants</li> <li>Human Reproduction</li> <li>Reproductive Health</li> </ol>
2. Magnetic Effect of Current and Magnetism	<ol> <li>Electrochemistry</li> <li>Surface Chemistry</li> </ol>	Module-2  1. Principles of Inheritance and
Module-3	Module-3(IC)	Variation  2. Molecular Basis of Inheritance
<ol> <li>Electromagnetic Induction</li> <li>Alternating Current</li> </ol>	<ol> <li>P-Block Elements</li> <li>Transition Elements</li> </ol>	3. Evolution
Module-4	(d & f block)  3. Co-ordination Compound	Module-3
<ol> <li>Geometrical Optics</li> <li>Wave Optics</li> </ol>	4. Metallurgy  Module-4(OC)	<ol> <li>Human Health and Disease</li> <li>Strategies for Enhancement in Food Production</li> </ol>
Module-5	1. HaloAlkanes & HaloArenes 2. Alcohol, Phenol & Ether	3. Microbes in Human Welfare
<ol> <li>Modern Physics</li> <li>Nuclear Physics</li> <li>Solids &amp; Semiconductor</li> </ol>	3. Aldehyde, Ketone & Carboxylic Acid	Module-4  1. Biotechnology: Principles and Processes
Devices 4. Electromagnetic Waves	Module-5(OC)  1. Nitrogen & Its Derivatives 2. Biomolecules & Polymers	<ul><li>2. Biotechnology and Its</li><li>Applications</li><li>3. Organisms and Populations</li></ul>

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**3.** Chemistry in Everyday Life