

Section (B) : Resonance Concepts, Conditions, Resonating Structures & Conjugation

B-1. Which of the following compounds have delocalized electrons ?





(S) CH₃CH₂NHCH=CHCH₃

B-2. Number of π electrons in resonance in the following structure is.



Section (C) : Stability of Resonating Structures and different species

C-1. In the following sets of resonating structure, label the major and minor contributors towards resonance hybrid.

C-2. Write the stability order of following resonating structures :



Section (D) : Mesomeric Effect

GOC-I

D-1. Arrange the following groups in the increasing order of +M:

(i)
$$-I$$
, $-CI$, $-F$, $-Br$ (ii) $-NH_2$, $-OH$, $-O^{\Theta}$

 $\label{eq:D-2.} \textbf{D-2.} \qquad \text{Arrange the following groups in the increasing order of } -M:$

D-3. Which of the following groups (attached with benzene ring) show +M effect?



Section (E) : Steric Inhibition of Resonance (SIR Effect)

E-1. Compare the SIR effect between orthochloro benzoic acid, orthobromobenzoic acid and orthoiodo benzoic acid.

Section (F) : Hyperconjugation

- **F-1.** Define hyperconjugation by taking an example of propene.
- **F-2.** In which molecules or ions hyperconjugation effect is observed and write the number of hyperconjugable hydrogen atoms.



Section (G) : Concept of Aromaticity

- G-1. What is aromaticity ?
- G-2. Classify the following as aromatic, antiaromatic and nonaromatic compounds.



G-3. Why cyclooctatetraene is nonplanar.

Section (H) : Applications of electronic effect

H-1. The correct decreasing order of electron density in aromatic ring of following compounds is :



(C) - OR

A-1. Inductive effect involves :

H-2.

- (A) Delocalisation of σ -electrons
- (C) Delocalisation of π -electrons
- (B) Partial displacement of σ -electrons

(D) Displacement of lone pair electrons.

- A-2. Select correct statement about I effect?
 - (A) I effect transfers electrons from one carbon atom to another.
 - (B) I effect is the polarisation of σ bond electrons.
 - (C) I effect creates net charge in the molecule.
 - (D) I effect is distance independent.
- A-3. Which of the following group shows +I-effect : (A) - Br(B)-COOH

"I was willing to accept what I couldn't change." - Dr. A. P. J. Abdul Kalam

(D) -COO-



Parishram

B-10. Which of the following pair is not pair of resonating structures?

GOC-I

(A)
$$(B) CH_2 = CH - O & CH_2 - CH = O$$

(C) $CH_2 = CH - O - CH_3 & CH_2 - CH = O - CH_3$
(D) $CH_3 - C = O & CH_3 - C = O$

B-11. Which of the following structures are resonance structures ?



B-12. Among the given sets, which represents the resonating structure ?

(A)
$$H-C \equiv \overset{\oplus}{N-O}$$
: and $H-O-C \equiv N$.
(B) $H-O = C = N$ and $H-O-C \equiv N$.
(C) $H-C \equiv \overset{\oplus}{N-O}$: and $H-C-N$.
(D) $H-O-C \equiv N$ and $H-N = C = O$.

B-13. In which of the following resonance is not possible? (A) $CH_2 = C = CH_2$ (B) $CH_2 = C = CH^{\odot}$ (C) $CH_2 = C = CH^{\odot}$ (D) $CH_2 = C = CH^{\bullet}$

Section (C) : Mesomeric Effect

- C-1. Which of the following group show +M effect? (A) -CN (B) -O-NO (C) $-CCI_3$ (D) -CHO
- C-2. Which of the following group show –M effect?

(A)
$$-CMe_{3}$$
 (B) $-O-S-O-R$ (C) $-NH-C-CH_{3}$ (D) $-S-O-R$

C-3. Which of the following group show +M and -I effect?

$$(A) -C -F \qquad (B) -C -OR \qquad (C) -O^{\Theta} \qquad (D) -OH$$

C-4. Which of the following group show +M > -I effect ?

(A)
$$-F$$
 (B) $\underset{-O-C-R}{\parallel}$ (C) $\underset{-O}{-C-R}$ (D) $-COOH$

C-6. +M and +I both effects are shown by : (A) $- \dddot{O}H$ (B) $- \dddot{N}HCH_3$ (C) $- O^{\ominus}$ (D) $- C (CH_3)_3$

C-7. The weakest +M group of the given species is : (A) $-OCH_3$ (B) -F (C) -I (D) $-N(CH_3)_2$

[&]quot;I was willing to accept what I couldn't change." - Dr. A. P. J. Abdul Kalam



Section (D) : Stability of Resonating Structures and different species

D-1. Which one of the following is least stable resonating structure ?

GOC-I

$$\begin{array}{c} O \\ \parallel \\ (A) \ \mathsf{NH}_2-\mathsf{C}-\mathsf{O}-\mathsf{CH}_3 \end{array} (B) \begin{array}{c} \Theta \\ \mathsf{NH}_2=\mathsf{C}-\mathsf{O}\mathsf{CH}_3 \end{array} (C) \ \mathsf{NH}_2-\overset{\mathsf{O}}{=}\mathsf{O}\mathsf{CH}_3 \end{array} (D) \ \mathsf{NH}_2-\overset{\mathsf{O}}{=}\mathsf{O}\mathsf{CH}_3$$

D-2. Which of the following resonating structure is the least contributing structure ?

$$(A) \xrightarrow{O^{\ominus}} (B) \xrightarrow{\ominus} (C) \xrightarrow{\oplus} (D) \xrightarrow{\oplus} (D)$$

D-3. HNCO (isocyanic acid) has following resonating structures :

$$\begin{array}{cccc} H-N=C=O & \longleftrightarrow & H-\overset{\odot}{N}-C=\overset{\oplus}{O} & \longleftrightarrow & H-\overset{\oplus}{N}\equiv C-\overset{\ominus}{O}\\ I & III & II$$

$$\begin{array}{c} (I) \\ (II) \\ (A) (IV) > (I) > (III) > (II) \\ (B) (II) > (IV) > (I) > (III) \\ (C) (III) > (IV) > (I) > (IV) > (IV) > (III) > (IV) > (II) > (IV) > (II) > (IV) > (III) > (III) > (IV) > ($$

D-5. Which is the most stable resonating structure ?





Section (E) : Steric Inhibition of Resonance (SIR Effect)

E-1. Maximum extent of steric inhibition of resonance can be expected in





E-2. Select the correct statement about this compound.



(A) All three C–N bond length are same.

- (B) C_1 –N and C_3 –N bonds length are same but shorter than C_5 –N bond length.
- (C) C_1 -N and C_5 -N bonds length are same but longer than C_3 -N bond length.
- (D) $C_1 N$ and $C_3 N$ bonds length are different but both are longer than $C_5 N$ bond length.

Section (F) : Hyperconjugation

F-1. In hyperconjugation there is overlap between : (A) p- and π -orbitals (B) 2π -orbitals (

(C) d-and- π -orbtial

(D) σ -and p - orbitals

Parishram

ÇH₃

F-2. Which of the following cannot exhibit hyperconjugation -

(A)
$$CH_3CH_2$$
 (B) $CH_3 CH_3 CH_3 CH_2$ (C) $CH_3CH = CH_2$ (D) $(CH_3)_3C - CH_2$

F-3. Which of the following alkenes will show maximum number of hyperconjugation forms ?

(A)
$$CH_2 = CH_2$$
 (B) $CH_3 - CH = CH_2$ (C) $CH_3 - CH_2 - CH_2 - CH_3$ (D) $CH_3 - CH_2 - CH_2$

F-4. Arrange the stability of following

GOC-I



F-5. Which one of the following has inductive, mesomeric and hyperconjugation effect ? (A) CH_3CI (B) $CH_3 - CH = CH_2$

(C)
$$CH_{3}CH = CH - C - CH_{3}$$
 (D) $CH_{2} = CH - CH = CH_{2}$

F-6. Which of the following group has the maximum hyperconjugation effect when attached to bezene ring ?(A) CH_3 -(B) CH_3CH_2 -(C) $(CH_3)_2CH$ -(D) $(CH_3)_3C$ -

Section (G) : Concept of Aromaticity

G-1. Which out of the following is aromatic hydrocarbon ?



G-2. Identify the aromatic compound ?



- G-3. Aromatic compounds burn with sooty flame because :
 - (A) They have a ring structure of carbon atoms.
 - (B) They have a relatively high percentage of hydrogen.
 - (C) They resist reaction with oxygen of air.
 - (D) They have a relatively high percentage of carbon.

Section (H) : Applications of electronic effect

H-1. The decreasing order of electron density on the ring is :





2. In which delocalisation of positive charge is possible ?

GOC-I



3. Decreasing order of potential energy of the following cations is :



4. Stability order of the following species ?



5. In which of the following first resonating structure is more stable than the second ?

6. Which of the following is **incorrect** for stability of structures.

7. Least contributing resonating structure of nitroethene is :

(A)
$$CH_{2} = CH - \dot{N} \bigcirc 0^{-}$$

(B) $\dot{C}H_{2} - \bar{C}H - \dot{N} \bigcirc 0^{-}$
(C) $\bar{C}H_{2} - \dot{C}H - \dot{N} \bigcirc 0^{-}$
(D) $\dot{C}H_{2} - CH = \dot{N} \bigcirc 0^{-}$

8. Which of the following statement is correct ?

(A) In the dianion (a), all the C–C bonds are of same length but C–O bonds are of different length

- (B) In the dianion, all C–C bonds are of same length and also all C–O bonds are of same lengths
- (C) In the dianion, all C–C bond lengths are not of same length
- (D) None of the above



10. Which of the following is correct about the following compound



(Naphthalene)

- (A) All the C-C bond length are same
- (B) $C_1 C_2$ bond length is shorter than $C_2 C_3$ bond length
- (C) $C_1 C_2$ bond length is greater than $C_2 C_3$ bond length
- (D) All the C-C bond length are equal to C-C bond length of benzene
- **11.** The correct order of +M effect of 'N' containing functional group on benzene ring, amongst the given compounds is



 $(A) I > II > IV > III \qquad (B) II > I > III > IV \qquad (C) I > II > III > IV \qquad (D) IV > III > II > II > I$

- **12.** In which case the σ -bond pair and π bond pair of electrons both are attracted in the same direction, (towards same atom.): (A) H₂C=CH-CI (B) CH₂-CH₂-NH₂ (C) H₂C=CH-CH=O (D) H₂C=CH-OCH₂
- **13.** The correct stability order of given resonating structures is :





14.

15. Select the correct order of heat of hydrogenation ?

GOC-I



- (A) The boron is sp² hybridized and the p-orbital contains an unshared pair of electron
- (B) The boron is sp² hybridized and a hybrid orbital contains an unshared pair of electron.
- (C) The boron in sp²hybridized and hybrid orbital is vacant
- (D) The boron is $sp^{\scriptscriptstyle 2}$ hybridized and the p–orbital is vacant
- 20. The correct order of electron density in aromatic ring of following compounds is :



GOC-I

Parishram





4. Identify the number of compounds in which positive charge will be delocalised ?





GOC-I



In how many of the following compounds Hyperconjugation effect is observed -6.



7.







8. Total number of moelcules which are antiaromatic?



Parishram

"I was willing to accept what I couldn't change." - Dr. A. P. J. Abdul Kalam



9. Find the number of carbon atoms including the given structure which can have negative change in resonating structures. (The structure with charge reperating are not accepted)



10. Observe the following compound and write the number of hydrogen atom involved in hyperconjugation?



11. Find the total number of positions where positive charge can be delocalized by true resonance



(Excluding the given position)

PART-III : MULTIPLE CHOICE QUESTION

- 1. Which statement is/are true about resonance?
 - (A) It decreases the energy of system.
 - (B) The hybridisation of atoms do not change due to resonance
 - (C) Resonance hybrid is more stable than any resonating structure.
 - (D) Resonanting structures can not be isolated at any temperature
- 2. Which of the following statement is incorrect about resonance ?
 - (A) The most stable structure explains all the characteristics of a species.
 - (B) All resonating structures remain in equilibrium.
 - (C) Resonance hybrid has maximum similarity with most stable resonating structure.
 - (D) Resonance hybrid is real.

"I was willing to accept what I couldn't change." - Dr. A. P. J. Abdul Kalam

3.

GOC-I

In which of the following pairs of compounds, will second structure have more contribution to resonance hybrid than first ?



4. In which of the following pairs of resonating structures first resonating structure is more stable than second?



5. In which of the following compounds delocalisation of electrons and shifting of electron in the same direction?

(B) CH₃-NH-NO₂

H (C) $CH_2=CH-CH=CH-NO_2$ (D) $H_2N-C=CH_2$ Which of the following groups cannot participate in resonance with benzene : (A) -COOH (B) COO^{\oplus} (C) $-\overset{\oplus}{N}H_3$ (D) Which of the following is/are correct : (A) is antiaromatic (B) is around the following is around the following is antiaromatic (B) is a round the following the following is a round the following the following is a round the following is a round the following t



Page No # 15

(D)-COCI

(A)

6.

7.



GOC-I





PART-IV : COMPREHENSION

Read the following passage carefully and answer the questions.

Hydrogenation of unsaturated hydrocarbons is an exothermic reaction. Due to hyperconjugation and resonance the stability of unsaturated hydrocarbons increases and the increase in stability is more due to resonance. Compound with same number of π -bonds and more stability has lower heat of hydrogenation. Heat of formation is defined as the energy evolved when a molecule is formed from its atoms. For isomers the more stable compound has higher heat of formation.

- 1.The correct heat of hydrogenation order is :
(p) 1, 3-Pentadiene
(r) 2, 3-Dimethyl-1,3-butadiene
(A) p > q > r > s(q) 1, 3-Butadiene
(s) Propadiene
(C) q > s > p > r(D) s > p > q > r > s(D) s > p > q > r > r
- 2. The order of heat of formation of the following molecules is :



9.

H₂Ċ–N=N

(||)

Parishram

(A) CH₂ at C-4

 $H_{0}C = \dot{N} = \dot{N}$

(I)

form any ion even in the presence of Ag⁺. Explain why ? Which will be the least stable resonating structure : [JEE-05(S), 3/84] (A) $CH_2 = CH - \overset{\textcircled{e}}{CH} - \overset{\textcircled{o}}{CH} - O - CH_3$ (B) $\overset{\textcircled{o}}{CH}_2 - \overset{\textcircled{o}}{CH} - CH = CH - OCH_3$ (C) $\overset{\Theta}{C}$ H₂ - CH = CH - CH = $\overset{\Theta}{O}$ - CH₃ (D) CH₂ = CH - $\overset{\Theta}{C}$ H - CH = $\overset{\Theta}{O}$ - CH₃ Among the following, the least stable resonating structure is : [JEE-07, 3/162] (D) (C) ⊕ Hyperconjugation involves overlap of the following orbitals : [JEE-08, 3/163] $(A) \sigma - \sigma$ (B) σ–p (C) p - p(D) $\pi - \pi$

PART-I : JEE PROBLEMS (PREVIOUS YEARS)

* Marked Questions may have more than one correct option.

- Write resonating structure of the compound H_3C CH_2 CH_2 OH1.
- 2. Explain why 7-bromo-1, 3, 5-cycloheptatriene exist as an ion while 5-Bromo-1, 3-cyclopentadiene does not [JEE 2004, 4/60]
- 3.
- 4.

5. The correct stability order of the following resonating structures is : 6. [JEE-09, 3/160]

H₂C–N≡N

(III)

(A) (I) > (II) > (IV) > (III)(B) (I) > (III) > (II) > (IV)(D) (III) > (I) > (IV) > (II)(C) (II) > (I) > (III) > (IV)

In the following carbocation; H/CH_a that is most likely to migrate to the positiviely charged carbon is 7.

> [JEE-09, 3/160] (B) H at C-4 $(C) CH_3 at C-2$ (D) H at C-2

 $H_{\lambda}\overline{C}-N=N$

(IV)

[JEE-03(S), 2/60]

TING THE TAR

Exercise #3

GOC-I

8. The total number of contributing structures showing hyperconjugation (involving C–H bonds) for the following carbocation is [JEE-11,4/180]



GOC-I

9.* Which of the following molecules, in pure form, is (are) unstable at room temperature ? [JEE-12, 4/136]



10. Find the number of resonance structures for N.



[IIT-2013]

PART-II: AIEEE PROBLEMS (PREVIOUS YEARS)

1. Picric acid is





[JEE MAIN-2002]

- 2. Which of the following species acts both as bronsted acid and base ? [JEE MAIN-2002] (A) NH₃ (B) OH_ (C) HSO⁻₄ (D) Both (A) and (C)
- $\begin{array}{lll} \textbf{3.} & \mbox{The correct order of increasing basic nature for the bases NH_3, CH_2NH_2 and (CH_3)_2NH is \\ (A) CH_3NH_2 < NH_3 < (CH_3)_2NH & (B) (CH_3)_2NH < NH_3 < CH_3NH_2 \\ (C) NH_3 < CH_3NH_2 < (CH_3)_2NH & (D) CH_3NH_2 < (CH_3)_2NH < NH_3 \\ \end{array}$
- 4. Arrange the carbanions, $(CH_3)_3 \overline{C}, \overline{C}CI_3, (CH_3)_2 \overline{C}H, C_6H_5\overline{C}H_2$, in order of their decreasing stability [JEE MAIN-2002]

(A)
$$\overline{C}CI_3 > C_6H_5\overline{C}H_2 > (CH_3)_2\overline{C}H > (CH_3)_3\overline{C}$$

(C) $C_6H_5\overline{C}H_2 > \overline{C}CI_3 > (CH_3)_3\overline{C} > (CH_3)_2\overline{C}H$

5. In the following benzyl/allyyl system

(R is alkyl group) The decreasing order of inductive effect is (A) $(CH_3)_3C \rightarrow (CH_3)_2CH \rightarrow CH_3CH_2 \rightarrow (CH_3)_2CH \rightarrow CH_3CH_2 \rightarrow (CH_3)_3CH \rightarrow CH_3CH_2 \rightarrow (CH_3)_3CH \rightarrow (CH_3)_3$

NO₂

OH

(B) CH_3 - CH_2 -> $(CH_3)_2CH$ -> $(CH_3)_3C$ -(D) None of these



Page No # 20

"I was willing to accept what I couldn't change." - Dr. A. P. J. Abdul Kalam

Page No # 21

GOC-I

Parishram

Parishram

Image: here in the following is more basic than aniline(3)
$$CH_3 - C_{H_3}^{H_3}$$
(4) $CH_3 - C_{H_3}^{\oplus}$ (5) CH_3^{O} (5) CH_3^{O} (5) CH_3^{O} (7) CH_3^{O}

(1)
$$FCH_2COOH > CH_3COOH > BrCH_2COOH > CICH_2COOH$$

(2) $BrCH_2COOH > CICH_2COOH > FCH_2COOH > CH_3COOH$

(3) $FCH_2COOH > CICH_2COOH > BrCH_2COOH > CH_3COOH$

(4) $CH_3COOH > BrCH_2COOH > CICH_2COOH > FCH_2COOH$

12. The stability of carbanions in the following:-

(a)
$$\mathbf{RC} \equiv \mathbf{C}^{\Theta}$$

Parishram

is in the order of:-

(1) (d) > (b) > (c) >(a) (2) (a) >(c) >(b) >(d) (3) (a) > (b) > (c) > (d) (4) (b) > (c) > (d) >(a)The state of hybridization of C_2 , C_3 , C_5 and C_6 of the hydrocarbon, 13.

$$\begin{array}{c} CH_{3} & CH_{3} \\ CH_{3}-C - CH_{3} - CH_{4} - CH_{3} - CH_{2} \\ CH_{3} \end{array}$$

is in the following sequence :-(1) sp, sp², sp³ and sp² (3) sp³, sp², sp² and sp (2) sp, sp³, sp² and sp³ (4) sp, sp², sp² and sp³

Which one of the following compounds has the most acidic nature ? 14.



GOC-I

7.

Which amongst the following is the most stable carbocation :-

[AIPMT-2005]

Parishram

6]

(c) $R_2C = CH$ (d) $R_3C - CH_2$

[AIPMT-2008]

[AIPMT-2010]

[AIPMT-2009]





have + M group.

- **E-1.** SIR effect increases with the size of ortho group. The order of SIR effect is o-iodo benzoic acid > o-bromo benzoic acid > o-chloro benzoic acid.
- F-1. It is delocalisation of sigma electron with p-orbital. It may take place in alkenes, alkynes, carbocations, free radicals, alkelyl benzene.

Necessary Condition : Presence of at least one hydrogen at saturated carbon which is α with respect to alkene, alkynes, carbocation, free radical, benzene nucleus.

Ż	Pa	ari	is	hr	aı	m
	TRANC	ATT MOVE	o rrroz	TT INTO	or ar one	10.000

(D) No hyperconjugation

(H) 9

(L) 10

GOC-I	

- F-2.
 (A) 0
 (B) 2
 (C) 0

 (E) 0
 (F) 0
 (G) 6

 (I) 3
 (J) 10
 (K) 0
- **G-1.** Those molecules are aromatic which have very high resonance energy. Only those molecules has sufficiently high amount of resonance energy to become aromatic which (a) are cyclic
 - (b) are planar
 - (c) contains (4n +2) number of π -electrons in ring.
 - (d) must have cyclic resonance between (4n + 2) number of π -electrons Where n = 0,1,2,3,4
- **G-2.** Aromatic : (a), (b), (c), (d), (e), (f), (h), (l) Antiaromatic : (g), (i), (j), (m) Nonaromatic : (k)
- G-3. Cyclooctatetraene is nonplanar to avoid its anti aromaticity and it becomes tub-shaped structure.

H-1. (III) > (IV) > (IV) > (I) +m group increases electron density and – m group decreases electron density in aromatic ring.

H-2. p > q > r

			PA	RT-	OBJE	CTIV	E QUES	TIC	DN
A-1.	(B)	А	-2.	(B)	A-3.	(D)	A	4.	(B)
A-6.	(A)	В	i-1 .	(A)	B-2.	(D)	B·	3.	(D)
B-5.	(A)	В	-6.	(A)	B-7.	(C)	B	·8.	(C)
B-10.	(A)	В	-11.	(B)	B-12.	(B)	B	13.	(A)
C-2.	(D)	C	-3.	(D)	C-4.	(B)	C-	5.	(A)
C-7.	(C)	D)-1.	(C)	D-2.	(B)	D	3.	(A)
D-5.	(B)	E	-1.	(C)	E-2.	(C)	F-	1.	(D)
F-3.	(B)	F	-4.	(C)	F-5.	(C)	F-	6.	(A)
G-2.	(B)	G	i-3.	(D)	H-1.	(A)	H	2.	(B)
H-4.	(A)								

PART-III : MATCH THE COLUMN

1. (A) - p,q,r,t ; (B) - p,q,r,t ; (C) - p,q,r,t ; (D) - p,q,s

	-(DUR EFFORTS INTO PERFORMANC DARDS KVPY OLYMPIAD
	Exerci	ise #	# 2		R		FOR C	HAL	LENG	ES	
~		P	ART-	:	OBJE		QUEST	ION			
1.	(C)	2.	(D)		3.	(A)	4.	(A)		5.	(C)
6.	(A)	7.	(C)		8.	(B)	9.	(B)		10.	(B)
11.	(C)	12.	(C)		13.	(D)	14.	(D)		15.	(B)
16.	(D)	17.	(A)		18.	(B)	19.	(D)		20.	(D)
	PAR	IT-II : 	SINGL	E AN	D DO	UBLE V	ALUE IN	TEGE	R TYP I		
1. 4.	5 (i, iii, iv, v, 5 (i, iii, v, vi,	vii) vii)		2. 5.	8 (ii, i 5 (i, ii	ii, iv, vi, vii, v , iv, vi, ix)	iii, ix, xi)	3. 6.	5 (i, ii, v 3 (B, C	, vi, ix) & G).	
7. 8	6 (i, iv, v, vi, 3	vii, x)									
0.	Aromatic –	a, c, g, h	, i, j, k.	;	Antia	romatic – b,	d, e ;	Nona	romatic – f		
9.	6			10.	9			11.	2		
		PART-		MUĽ	TIPL	E CHO	ICE OU	EST	ION		
1.	(ABCD)	2.	(AB)		3.	(ABC)	4.	(BC)		5	(BC)
6	(C)	7.	(BD)		8.	(AD)	9.	(ABD)	1		
			PAR	T-IV	: C(MPRE	HENSIC				
1.	(B)	2.	(C)								
	Exerci	ise <i>‡</i>	# 3		i				AKG	EL	
		PAR	T-I : J	EE PF	ROBLE	MS (PRE	VIOUS Y	EARS]		
1	CH3	ČH ₂									
	∥ ⊕ОН										
2.	7-bromo-1,	3, 5-cyclol	heptatrier	ne on ior	nisation (gives tropyliu	m ion	which is	aromatic	& highly	y stable
									Ð		
	but ionisatio	n of 5-bron	no-1, 3-cy	vclopenta	adiene gi	ves 1, 3-cyclo	opentadienyl	cation (which	is anti a	aromati
	& UNSTADIE.	(non exist	lent)								

(GOC-I	<u> </u>							Shram FFORTS INTO PERFORMANCE DS KVPY OLYMPIADS
3.	(A)	4.	(A)	5.	(B)	6.	(B)	7.	(D)
8.	6	9.*	(BC)	10.	9				
		PART-I	I : AIEEE PI	ROBLE	MS (PREVI	OUS	YEARS)		
1.	(C)	2.	(D)	3.	(D)	4.	(A)	5.	(A)
6.	(D)	7.	(C)	8.	(C)	9.	(D)	10.	(C)
11.	(D)	12.	(C)	13.	(A)	14.	(B)	15.	(D)
16.	(B)	17.	(C)	18.	(A)	19.	(A)	20.	(B)
21.	(A)	22.	(C)	23.	(B)	24.	(D)	25.	(D)
26.	(C)	27.	(D)	28.	(B)	29.	(C)	30.	(C)
31.	(A)		. ,				. ,		

PART-III : NEET PROBLEMS (PREVIOUS YEARS)

Que.	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
Ans.	1	4	3	4	1	2	3	1	2	4	3	3	2	3	2
Que.	16	17	18	19	20	21									
Ans.	4	2	3	4	1	3									

Parishram