

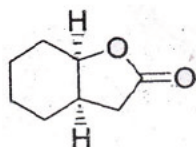
SOLVED PAPER : CSIR-UGC-NET/JRF- Dec. 2011**CHEMICAL SCIENCES BOOKLET - [C]****PART -B**

21. Identify which of the following operators is not hermitian ?
(a) $\frac{\hbar}{i} \frac{d}{dx}$ (b) $i \frac{d^2}{dx^2}$ (c) $\frac{d^2}{dx^2}$ (d) x^2
22. The term symbol for the ground state of nitrogen atoms is
(a) 3P_0 (b) $^4P_{3/2}$ (c) 1P_1 (d) $^4S_{3/2}$
23. P_A and P_B denote the populations of two energy states E_A and E_B , and $E_A > E_B$. The correct statement when the temperature $T_1 > T_2$ is
(a) $P_A(T_1) > P_B(T_1)$, $P_A(T_2) < P_B(T_2)$ and $(P_A/P_B)_{T_1} > (P_A/P_B)_{T_2}$
(b) $P_A(T_1) < P_B(T_1)$, $P_A(T_2) > P_B(T_2)$ and $(P_A/P_B)_{T_1} < (P_A/P_B)_{T_2}$
(c) $P_A(T_1) < P_B(T_1)$, $P_A(T_2) < P_B(T_2)$ and $(P_A/P_B)_{T_1} > (P_A/P_B)_{T_2}$
(d) $P_A(T_1) < P_B(T_1)$, $P_A(T_2) < P_B(T_2)$ and $(P_A/P_B)_{T_1} < (P_A/P_B)_{T_2}$
24. The uncertainty in the NMR frequency of a compound in liquid state (relaxation time = 1s) is 0.1 Hz. The uncertainty in the frequency (in Hz) of same compound in solid state (relaxation time = 10^{-4} s) is
(a) 10^{-4} (b) 100 (c) 1000 (d) 10^{-3}
25. Which one of the following conductometric titrations will show a linear increase of the conductance with volume of the titrant added up to the break point and an almost constant conductance afterwards.
(a) A strong acid with a strong base (b) A strong acid with a weak base
(c) A weak acid with a strong base (d) A weak acid a weak base.
26. Flocculation value of K_2SO_4 is much less than that of KBr for Sol A. Flocculation value of $CaCl_2$ is much less than that of NaCl for Sol B. Which of the following statements is correct ?
(a) Sol A is negatively charged and Sol B is positively charged
(b) Both the sols are negatively charged.
(c) Sol A is positively charged and sol B is negatively charged
(d) Both the sols are positively charged.
27. For a system of constant composition, the pressure (P) is given by.
(a) $-\left(\frac{\partial U}{\partial S}\right)_V$ (b) $-\left(\frac{\partial U}{\partial V}\right)_S$ (c) $\left(\frac{\partial V}{\partial S}\right)_T$ (d) $\left(\frac{\partial U}{\partial V}\right)_T$
28. The value of d_{111} in a cubic crystal is 325.6 pm. The value of d_{333} is
(a) 325.6 pm (b) 976.8 pm (c) 108.5 pm (d) 625.6 pm
29. The symmetry point group of ethane in its staggered conformation is
(a) C_{3v} (b) D_{3d} (c) D_{3h} (d) S_6

30. For the reaction $\text{C}_2\text{H}_4(\text{g}) + 3\text{O}_2(\text{g}) \rightarrow 2\text{CO}_2(\text{g}) + 2\text{H}_2\text{O}(\ell)$, the value of $\Delta H - \Delta U$ (in kJ) at 300 K and 1 bar is
 (a) -5.0 (b) 0.0 (c) 2.5 (d) 5.0
31. The sodium D lines are due to ${}^2\text{P}_{3/2} \rightarrow {}^2\text{S}_{1/2}(\Delta E_1)$ and ${}^2\text{P}_{1/2} \rightarrow {}^2\text{S}_{1/2}(\Delta E_2)$ transitions. The splitting due to spin-orbit coupling in ${}^2\text{P}$ state of the sodium atom is
 (a) $\Delta E_2 + \Delta E_1$ (b) $\Delta E_2 - \Delta E_1$ (c) $\frac{\Delta E_2 + \Delta E_1}{2}$ (d) $\frac{\Delta E_2 - \Delta E_1}{2}$
32. The rate constant of a unimolecular reaction was $2.66 \times 10^{-3} \text{ s}^{-1}$ and $2.2 \times 10^{-1} \text{ s}^{-1}$ at $T=120\text{K}$ and 360 K respectively. The rate constant (in s^{-1} units) at 240 K would be
 (a) 2.4×10^{-2} (b) 2.4×10^{-1} (c) 4.8×10^{-2} (d) 1.8×10^{-3}
33. For a potentiometric titration, in the curve of emf (E) vs volume (V) of the titrant added, the equivalence point is indicated by
 (a) $|dE/dV| = 0, |d^2E/dV^2| = 0$ (b) $|dE/dV| = 0, |d^2E/dV^2| > 0$
 (c) $|dE/dV| > 0, |d^2E/dV^2| = 0$ (d) $|dE/dV| > 0, |d^2E/dV^2| > 0$
34. The osmotic pressure (π) of a polymer sample at different concentrations (c) was measured at T(K). A plot of (π/c) versus c gave a straight line with slope (m) and intercept (c'). The number average molecular weight of the polymer is (R = gas constant).
 (a) $\frac{Rt}{c'}$ (b) $\frac{c'}{RT}$ (c) RT (d) mRT
35. The concentration of a reactant undergoing decomposition was 0.1, 0.08 and 0.067 mol L^{-1} after 1.0, 2.0 and 3.0 hr respectively. The order of the reaction is
 (a) 0 (b) 1 (c) 2 (d) 3
36. A particle is constrained in a one dimensional box of length 2a with potential $V(x) = \infty; x < -a, x > a$ and $V(x) = 0; -a \leq x \leq a$. Energy difference between levels $n = 3$ and $n = 2$ is
 (a) $\frac{5h^2}{8ma^2}$ (b) $\frac{9h^2}{8ma^2}$ (c) $\frac{9h^2}{32ma^2}$ (d) $\frac{5h^2}{32ma^2}$
37. In the ${}^{19}\text{F}$ NMR spectrum of PF_5 , the number of signals and multiplicity, at room temperature are
 (a) one, singlet (b) one, doublet (c) two, doublet (d) two singlet
38. The correct statement regarding closo- $\{\text{B}_n\text{H}_n\}$ species is :
 (a) it always has -2 charge
 (b) it always has +2 charge
 (c) it is a neutral species
 (d) it is more reactive than nido arachno-, and hypo-boranes
39. Lewis acidity of $\text{BCl}_3, \text{BPh}_3$ and BMe_3 with respect to pyridine follows the order
 (a) $\text{BCl}_3 > \text{BPh}_3 > \text{BMe}_3$ (b) $\text{BMe}_3 > \text{BPh}_3 > \text{BCl}_3$
 (c) $\text{BPh}_3 > \text{BMe}_3 > \text{BCl}_3$ (d) $\text{BCl}_3 > \text{BMe}_3 > \text{BPh}_3$

40. Superoxide dismutase contains the metal ions
 (a) Zn (II) and Ni(II) (b) Cu(II) and Zn(II)
 (c) Ni (II) and Co(III) (d) Cu (II) and Fe(III)
41. The number of antibonding electrons in NO and CO according to MO theory are respectively.
 (a) 1, 0 (b) 2, 2 (c) 3, 2 (d) 2, 3
42. The correct combination of metal, number of carbonyl ligands and the charge for a metal carbonyl complex $[M(CO)_x]^{z-}$ that satisfies the 18 electron rule is
 (a) M = Ti, x = 6, z = 1 (b) M = V, x = 6, z = 1
 (c) M = Co, x = 4, z = 2 (d) M = Mo, x = 5, z = 1
43. Among the following pairs
 (1) oxygen-sulfur (2) nitrogen-phosphorus
 (3) phosphorus arsenic (4) chlorine-iodine
 Those in which the first ionization energies differ by more than 300 kJ mole^{-1} are :
 (a) (1) and (3) only (b) (1) and (2) only (c) (2) and (3) only (d) (3) and (4) only
44. The stable cyclopentadienyl complex of beryllium is
 (a) $[Be(\eta^2-C_5H_5)_2]$ (b) $[Be(\eta^2-C_5H_5)(\eta^3-C_5H_5)]$
 (c) $[Be(\eta^1-C_5H_5)(\eta^3-C_5H_5)]$ (d) $[Be(\eta^1-C_5H_5)(\eta^5-C_5H_5)]$
45. The reaction between NH_4Br and Na metal in liquid ammonia (solvent) results in the products
 (a) NaBr, HBr (b) NaBr, H_2 (c) H_2 , HBr (d) HBr, H_2
46. The material that exhibits the highest electrical conductivity among the following sulfur-nitrogen compounds is
 (a) S_4N_4 (b) S_7NH (c) S_2N_2 (d) $(SN)_x$
47. Uranium fluorides co-precipitate with
 (a) CaF_2 (b) AgF (c) LiF (d) MgF_2
48. The acid-base indicator (HIn) shows a colour change at pH 6.40 when 20% of it is ionized. The dissociation constant of the indicator is
 (a) 9.95×10^{-8} (b) 3.95×10^{-6} (c) 4.5×10^{-8} (d) 6.0×10^{-8}
49. The actual magnetic moment shows a large deviation from the spin-only formula in the case of
 (a) Ti^{3+} (b) V^{3+} (c) Gd^{3+} (d) Sm^{3+}
50. The complex that absorbs light of shortest wavelength is
 (a) $[CoF_6]^{3-}$ (b) $[Co(H_2O)_6]^{3+}$ (c) $[Co(NH_3)_6]^{3+}$ (d) $[Co(OX)_3]^{3-}$ ($OX = C_2O_4^{2-}$)
51. Two α particles having speeds S_1 and S_2 have kinetic energies 1 and 2 MeV respectively; the relationship between S_1 and S_2 is :
 (a) $S_1 = 2S_2$ (b) $S_2 = 2S_1$ (c) $S_2 = \sqrt{2} S_1$ (d) $S_1 = \sqrt{2} S_2$

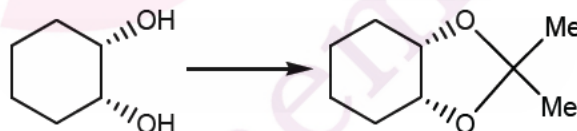
52. Green coloured $\text{Ni}(\text{PPh}_2\text{Et})_2 \text{Br}_2$, has a magnetic moment of 3.20 B.M. The geometry and the number of isomers possible for the complex respectively, are
 (a) square planar and one (b) tetrahedral and one
 (c) square planar and two (d) tetrahedral and two
53. The chemiluminescence method for determining NO in environmental samples is based on formation of NO_2^* (excited) which is generally generated by reacting NO with
 (a) O_2 (b) O_2^- (c) O_3 (d) O_2^{2-}
54. In the IR spectrum, carbonyl absorption band for the following compound appears at



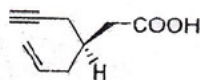
- (a) 1810 cm^{-1} (b) 1770 cm^{-1} (c) 1730 cm^{-1} (d) 1690 cm^{-1}
55. Among the following compounds, the formyl anion equivalent is
 (a) acetylene (b) nitromethane (c) ethyl chloroformate (d) 1, 4-dithiane
56. In the following concerted reaction, the product is formed by a



- (a) 6π -disrotatory electrocyclicalisation (b) 4π -disrotatory electrocyclicalisation
 (c) 6π -conrotatory electrocyclicalisation (d) 4π -conrotatory electrocyclicalisation.
57. A suitable reagent combination for carrying out the following conversion is

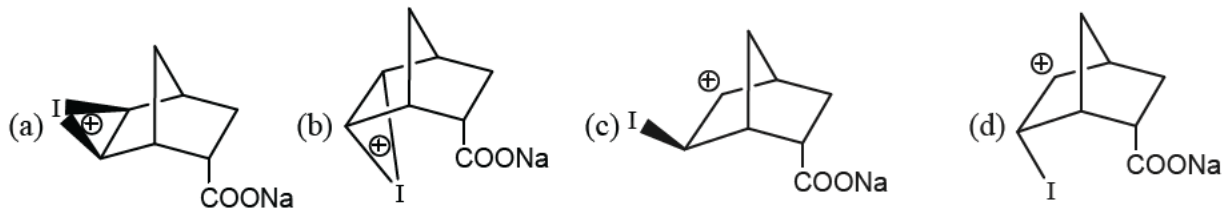
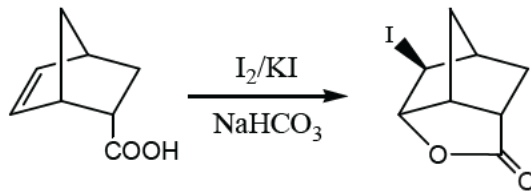


- (a) trimethyl orthoacetate and p-toluenesulfonic acid
 (b) trimethyl ortho acetate and sodium hydroxide
 (c) 2-methoxypropene and p-toluenesulfonic acid
 (d) 2-methoxypropene and sodium hydroxide
58. The IUPAC name of the following compound is



- (a) (R)-3- (prop-2-enyl) hex-5-ynoic acid (b) (S)-3- (prop-2-enyl) hex-5-ynoic acid
 (c) (R)-3- (prop-2-enyl) hex-5-enoic acid (d) (S)-3- (prop-2-ynyl) hex-5- enoic acid
59. In the mass spectrum of dodecahedrane ($\text{C}_{20}\text{H}_{20}$), approximate ratio of the peaks at m/z 260 and 261 is:
 (a) 1 : 1 (b) 5 : 1 (c) 10 : 1 (d) 20 : 1

60. The reaction given below proceeds through



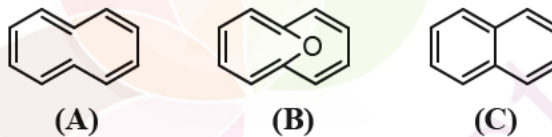
61. Among the following drugs, the anticancer agents is:

- (a) captopril (b) chloroquine (c) camptothecin (d) ranitidine

62. The reaction that involves the formation of both C-C and C-O bonds is

- (a) Diels-Alder reaction (b) Darzen's glycidic ester condensation
(c) aldol reaction (d) Beckmann rearrangement

63. Among A-C, the aromatic compounds are



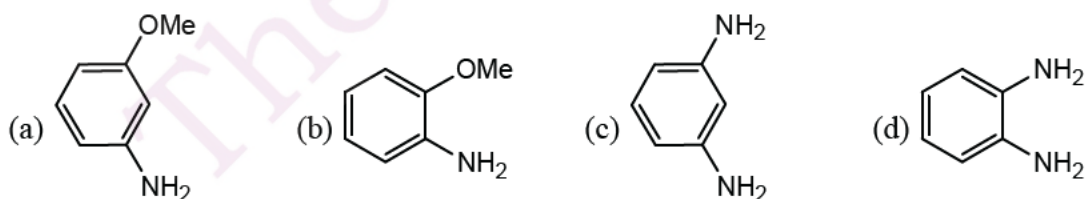
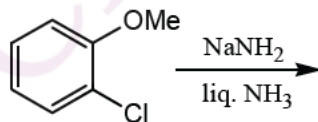
- (a) A, B and C (b) A and B only (c) B and C only (d) A and C only

64. In the following Markownikov addition reaction, the products A and B are

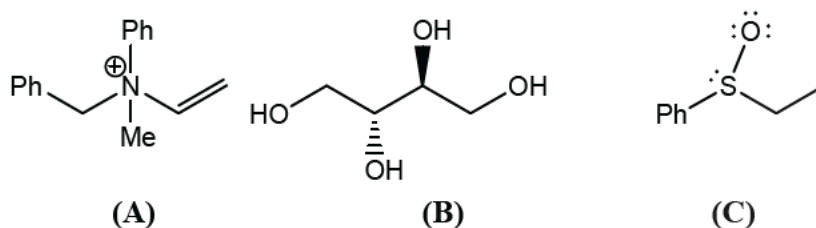


- (a) homomers (b) enantiomers (c) diastereomers (d) regioisomers

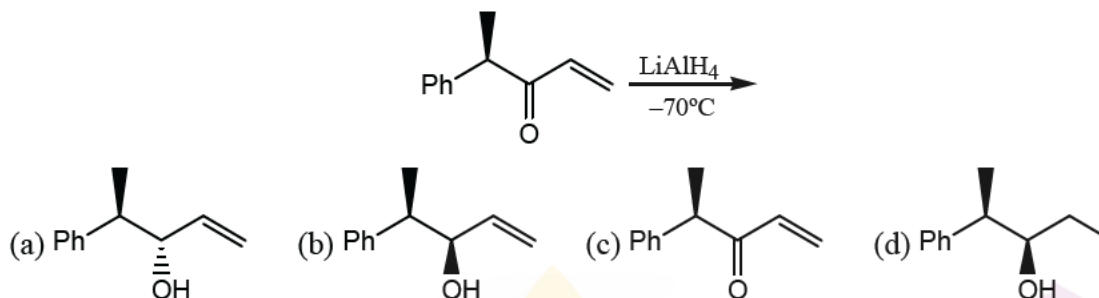
65. The major product formed in the following reaction is



66. Among A-C, the compounds which can exhibit optical activity are



- (a) A, B and C (b) A and B only (c) A and C only (d) B and C only
67. The major product formed in the following reaction is

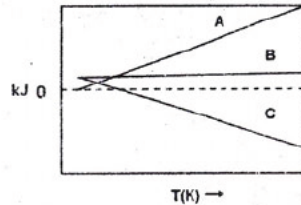


68. An organic compound (MF: $C_8H_{10}O$) exhibited the following 1H NMR spectral data : δ 2.5 (3 H,s), 3.8 (3H, s), 6.8 (2 H, d, J=8 Hz), 7.2 (2 H, d, J=8 Hz) ppm. The compound among the choices, is
- (a) 4-ethylphenol (b) 2-ethylphenol (c) 4-methylanisole (d) 4-methylbenzyl alcohol
69. With respect to electrophilic aromatic substitution, reactivity order of pyrrole, pyridine and indole is
- (a) indole > pyrrole > pyridine (b) pyrrole > pyridine > indole
 (c) pyrrole > indole > pyridine (d) indole > pyridine > pyrrole
70. The most appropriate reagent suitable for the conversion of 2-optyne into trans-2-octene is
- (a) zinc and acetic acid (b) 10% Pd/C
 (c) lithium in liquid ammonia (d) hydrazine hydrate

PART C

71. Consider a n-type semiconductor whose $E_v = 0$, $E_c = 2.0\text{eV}$ and $E_d = 1.98\text{eV}$. The correct statement among the following is
- (a) $E_f = 1\text{eV}$ and is independent of T
 (b) $E_f = 1.99\text{eV}$ and remains independent of T
 (c) $E_f = 1.99\text{eV}$ and increases towards 2.0 eV with increase of T
 (d) $E_f = 1.99\text{eV}$ and decreases with increase of T.
72. Reaction of $Fe(CO)_5$ with OH^- leads to complex A which on oxidation with MnO_2 gives B. Compounds A and B respectively are
- (a) $[HFe(CO)_4]^-$ and $Fe_3(CO)_{12}$ (b) $[Fe(CO)_5(OH)]^-$ and $Fe_2(CO)_9$
 (c) $[Fe(CO)_4]^{2-}$ and $Mn_2(CO)_{10}$ (d) $[HFe(CO)_4]^-$ and Fe_2O_3

73. For the reaction $\text{H}_2\text{O}(\text{g}) + \text{C}(\text{graphite}) \rightleftharpoons \text{CO}(\text{g}) + \text{H}_2\text{O}(\text{g})$, the variation of energy parameter ΔG , ΔH and $T\Delta S$ of the reaction over a large temperature range is shown below. The correct identification of the curves is given by

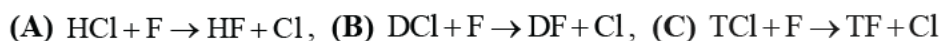


- (a) $A \rightarrow \Delta G^\circ$, $B \rightarrow \Delta H^\circ$, $C \rightarrow T\Delta S^\circ$ (b) $A \rightarrow \Delta H^\circ$, $B \rightarrow \Delta G^\circ$, $C \rightarrow T\Delta S^\circ$
 (c) $A \rightarrow \Delta G^\circ$, $B \rightarrow T\Delta S^\circ$, $C \rightarrow \Delta H^\circ$ (d) $A \rightarrow T\Delta S^\circ$, $B \rightarrow \Delta H^\circ$, $C \rightarrow \Delta G^\circ$
74. A Sodalite cage in zeolites is
 (a) a truncated tetrahedron (b) an icosahedron
 (c) a truncated octahedron (d) a dodecahedron
75. Two moles of a nonvolatile solute is dissolved in 48 mol of water and the resultant solution has a vapour pressure of 0.0392 bar at 300 K. If the vapour pressure of pure water at 300 K is 0.0400 bar, the activity coefficient of water in the solution is:
 (a) 0.96 (b) 0.98 (c) 1.00 (d) 1.02
76. The final product (s) of the reaction $\text{P}(\text{OR})_3 + \text{R}'\text{X}$ is/are
 (a) $\text{R}'\text{PO}(\text{OR})_2$ and RX (b) $[\text{R}'\text{PO}(\text{OR})_2]\text{X}$
 (c) $[\text{R}'\text{RPO}_2(\text{OR})]\text{X}$ (d) ROR' and $\text{P}(\text{OR})_2\text{X}$
77. 1 mol of CO_2 , 1 mol of N_2 and 2 mol of O_2 were mixed at 300 K. The entropy of mixing is
 (a) $6R \ln 2$ (b) $8R \ln 2$ (c) $\frac{8R \ln 2}{300}$ (d) $16R \ln 2$
78. For the eigenstates of the hydrogen atom, which of the following relations between the expectation value of kinetic energy (T) and potential (V) holds true?
 (a) $\langle T \rangle = \langle V \rangle$ (b) $2\langle T \rangle = -\langle V \rangle$ (c) $2\langle T \rangle = \langle V \rangle$ (d) $\langle T \rangle = -2\langle V \rangle$
79. For the liquid \rightleftharpoons vapour equilibrium of a substance $\frac{dP}{dT}$ at 1 bar and 400 K is $8 \times 10^{-3} \text{ bar K}^{-1}$. If the molar volume in the vapour form is 200 L mol^{-1} and the molar volume in the liquid form is negligible, the molar enthalpy of vapourisation is ($1.0 \text{ bar L} = 100 \text{ J}$)
 (a) 640 kJ mol^{-1} (b) 100 kJ mol^{-1} (c) 80 kJ mol^{-1} (d) 64 kJ mol^{-1}

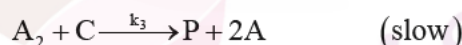
80. The correct order of acidity among the following species is
- (a) $[\text{Na}(\text{H}_2\text{O})_6]^+ > [\text{Ni}(\text{H}_2\text{O})_6]^{2+} > [\text{Mn}(\text{H}_2\text{O})_6]^{2+} > [\text{Sc}(\text{H}_2\text{O})_6]^{3+}$
 (b) $[\text{Sc}(\text{H}_2\text{O})_6]^{3+} > [\text{Ni}(\text{H}_2\text{O})_6]^{2+} > [\text{Mn}(\text{H}_2\text{O})_6]^{2+} > [\text{Na}(\text{H}_2\text{O})_6]^+$
 (c) $[\text{Mn}(\text{H}_2\text{O})_6]^{2+} > [\text{Ni}(\text{H}_2\text{O})_6]^{2+} > [\text{Sc}(\text{H}_2\text{O})_6]^{3+} > [\text{Na}(\text{H}_2\text{O})_6]^+$
 (d) $[\text{Sc}(\text{H}_2\text{O})_6]^{3+} > [\text{Na}(\text{H}_2\text{O})_6]^+ > [\text{Ni}(\text{H}_2\text{O})_6]^{2+} > [\text{Mn}(\text{H}_2\text{O})_6]^{2+}$
81. The Langmuir adsorption isotherm is given by $\theta = \frac{Kp}{1+Kp}$, where P is the pressure of the adsorbate gas. The Langmuir adsorption isotherm for a diatomic gas A_2 undergoing dissociative adsorption is:
- (a) $\theta = \frac{Kp}{1+Kp}$ (b) $\theta = \frac{2Kp}{1+2Kp}$ (c) $\theta = \frac{(Kp)^2}{1+(Kp)^2}$ (d) $\theta = \frac{(Kp)^{1/2}}{1+(Kp)^{1/2}}$
82. The standard electrode potentials (E^0) of $\text{Fe}^{3+}/\text{Fe}^{2+}$ and Fe^{2+}/Fe electrodes are +0.77V and -0.44 V respectively at 300 K. The E^0 of Fe^{3+}/Fe electrode at the same temperature is
- (a) 1.21 V (b) 0.33 V (c) -0.11V (d) -0.04 V
83. Which of the following is true for the radial part of the hydrogen atom wavefunctions $R_{nl}(r)$ (n principal quantum number) and the nodes associated with them?
- (a) The radial part of only s function is non-zero at the origin and has (n – 1) nodes.
 (b) The radial part of s function is zero at the origin and has n number of nodes.
 (c) All radial functions have values of zero at the origin and have (n – 1) nodes.
 (d) The radial parts of all s functions are zero at the origin and have no nodes.
84. For non-degenerate perturbation theory for ground state, with $E_0^{(0)}$ as zeroth order energy, $E_0^{(1)}$ as the first-order perturbation correction and E_0 as the exact energy, which of the following is true?
- (a) $(E_0^{(0)} + E_0^{(1)})$ is always equal to E_0 (b) $(E_0^{(0)} + E_0^{(1)}) \leq E_0$
 (c) $(E_0^{(0)} + E_0^{(1)}) \geq E_0$ (d) $E_0^{(0)} \leq (E_0 + E_0^{(1)})$
85. Observe the following electronic transition of a diatomic molecule.
- (a) ${}^1\Sigma_g^+ \rightarrow {}^3\Sigma_g^+$ (b) ${}^1\Sigma_u^+ \rightarrow {}^1\Sigma_g^+$ (c) ${}^1\Delta_u \rightarrow {}^1\Sigma_g^+$ (d) ${}^1\Pi_g \rightarrow {}^1\Sigma_u^+$
- The allowed transitions are
- (a) (A) and (C) only (b) (B) and (D) only
 (c) (A), (B) and (C) only (d) (A), (C) and (D) only

86. An excited triplet state wave function of hydrogen molecule with the electronic configuration $\sigma_g^1 \sigma_u^1$ has the following space part
- (a) $\sigma_g(1)\sigma_u(2)$ (b) $\sigma_g(1)\sigma_u(2) + \sigma_u(1)\sigma_g(2)$
 (c) $\sigma_g(1)\sigma_u(2) - \sigma_u(1)\sigma_g(2)$ (d) $\sigma_g(1)\sigma_g(2) + \sigma_u(1)\sigma_u(2)$
87. The NMR spectrum of AX_3 exhibits lines at $\delta = 2.1$ and 2.3 ppm (for X type protons) and $\delta = 4.1, 4.3, 4.5$ and 4.7 ppm (for A type protons), measured from TMS with an instrument operating at 100 MHz. The chemical shift (in ppm) of A and X protons and coupling constant (in Hz) are respectively.
- (a) 4.4, 2.2 and 20 (2) 2.2, 4.4 and 10 (c) 2.2, 4.4 and 5 (d) 4.3, 2.1 and 20
88. The character table of the C_{2v} point group is given below:
- | C_{2v} | E | C_2 | σ_v | σ'_v |
|----------|---|-------|------------|-------------|
| A_1 | 1 | 1 | 1 | 1 |
| A_2 | 1 | 1 | -1 | -1 |
| B_1 | 1 | -1 | 1 | -1 |
| B_2 | 1 | -1 | -1 | 1 |
- The two functions $\phi_1 = p_1 + 2p_2 + 2p_3 + p_4$ and $\phi_2 = 2p_1 - p_2 - p_3 + 2p_4$ (where p_k is the p-orbital on the k^{th} atom of cis-butadiene and σ_v is the molecular plane) belong to
- (a) A_1 and A_2 respectively (b) Both A_2
 (c) Both B_2 (d) B_1 and B_2 respectively.
89. If θ_r denotes the characteristic temperature of rotation then the magnitude of $[\theta_r(H_2)\theta_r(D_2)] / [\theta_r(HD)]^2$ (assume the bond lengths to be the same for all the molecules) is
- (a) 2/3 (b) 3/2 (c) 8/9 (d) 9/8
90. The overall reaction for the passage of 1.0 faraday of charge in the following cell $Ag(s) - AgCl(s) | KCl(a_1) | KCl(a_2) | AgCl(s) - Ag(s)$ is given by (t denotes the transport numbers)
- (a) $t_+ KCl(a_1) \rightarrow t_+ KCl(a_2)$ (b) $t_+ KCl(a_2) \rightarrow t_+ KCl(a_1)$
 (c) $t_- KCl(a_1) \rightarrow t_- KCl(a_2)$ (d) $t_- KCl(a_2) \rightarrow t_- KCl(a_1)$
91. A system consisting of 4 identical and distinguishable particles, each possessing three available states of 1, 2 and 3 units, has 10 units of energy. The number of ways, W , in which these conditions are satisfied is
- (a) 2 (b) 4 (c) 6 (d) 10
92. The molar conductivities at infinite dilution Λ_m^0 for $Na_2SO_4, K_2SO_4, KCl, HCl$ and $HCOONa$ at 300 K are 260, 308, 150, 426 and 105 $S\text{ cm}^{-1}\text{ mol}^{-2}$ respectively. Hence Λ_m^0 for formic acid in the same unit and at the same temperature is
- (a) 381 (b) 405 (c) 429 (d) 531

93. If the displacement vectors of all atoms in cis-butadiene are taken as the basis vectors the characters of the reducible representation of E , C_2 , σ_v (molecular plane) and σ_v' are
 (a) 30, 10, 30, 0 (b) 30, 0, 10, 0 (c) 30, 20, 0, 0 (d) 30, 0, 20, 0
94. In least square fitting of a data set $\{X_i, Y_i\}$ to the equation $Y = AX$, the regression coefficient (A) is estimated by
 (a) $\sum Y_i^2 / \sum X_i^2$ (b) $\sum X_i Y_i / \sum X_i^2$ (c) $\sum X_i Y_i / \sum Y_i^2$ (d) $\sum X_i^2 / \sum Y_i^2$
95. At any temperature for the following reaction (D and T are deuterium and tritium respectively) correct statement is:

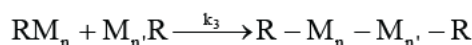
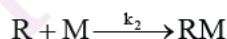


- (a) (A) is fastest (b) (B) is fastest (c) (C) is fastest
 (d) All the above reactions have the same rate constant.
96. An example of a relaxation method of measuring rates is:
 (a) Spectroscopic monitoring of product concentration.
 (b) Stopped flow technique
 (c) Temperature jump experiments.
 (d) Measurement of spectral line widths.
97. The overall rate of the following complex reaction,



by steady state approximation would be

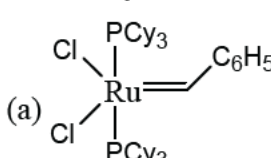
- (a) $K_1 K_2 k_3 [A]^3 [B]$ (b) $K_2 K_1 k_3 [A][B]^3$ (c) $K_1 K_2 k_3 [A][B]^2$ (d) $K_1 K_2 k_3 [A][B]$
98. The vibrational energy levels, $\nu'' = 0$ and $\nu' = 1$ of a diatomic molecule are separated by 2143 cm^{-1} . Its anharmonicity ($\omega_e x_e$) is 14 cm^{-1} . The values of ω_e (in cm^{-1}) and first overtone (cm^{-1}) of this molecule are respectively.
 (a) 2143 and 4286 (b) 2157 and 4286 (c) 2157 and 4314 (d) 2171 and 4258
99. The addition polymerization of M (monomer) involves the following stages:
 (I = initiator, R = free radical)



The rate constant for free radical formation is $2 \times 10^{-3} \text{ s}^{-1}$. The initial concentration of initiator is $10^{-3} \text{ mol dm}^{-3}$. The overall rate of the reaction is $4 \times 10^{-3} \text{ mol dm}^{-3} \text{ s}^{-1}$. Assuming steady state approximation for free radical, the kinetic chain length is:

- (a) 2000 (b) 8×10^9 (c) 20 (d) 200

100. The electronic spectrum of $[\text{CrF}_6]^{3-}$ shows three bands at $14,900\text{ cm}^{-1}$, $22,400\text{ cm}^{-1}$ and $34,800\text{ cm}^{-1}$. The value of Δ_0 in this case is:
- (a) $5,500\text{ cm}^{-1}$ (b) $14,900\text{ cm}^{-1}$ (c) $22,400\text{ cm}^{-1}$ (d) $34,800\text{ cm}^{-1}$
101. Among the following pairs, those in which both species have similar structures are:
- (A) N_3^- , XeF_2 (B) $[\text{ICl}_4]^-$, $[\text{PtCl}_4]^{2-}$ (C) $[\text{ClF}_2]^+$, $[\text{ICl}_2]^-$ (D) XeO_3 , SO_3
- (a) (A) and (B) only (b) (A) and (C) only
(c) (A), (B) and (C) only (d) (B), (C) and (D) only
102. The number of metal-metal bonds in the dimers, $[\text{CpFe}(\text{CO})(\text{NO})]_2$ and $[\text{CpMo}(\text{CO})_3]_2$ respectively, are
- (a) two and two (b) two and three (c) one and two (d) zero and one
103. The reduction of nitrogen to ammonia, carried out by the enzyme nitrogenase, needs,
- (a) 2 electrons (b) 4 electrons (c) 6 electrons (d) 8 electrons
104. In the titration of 50 mL of 0.1 M HCl with 0.1 M NaOH using methyl orange as an indicator, the end point (color change) occurs as pH reaches 4.0. The titration error is:
- (a) -0.2% (b) -84.7% (c) $+0.2\%$ (d) $+84.2\%$
105. The styx code of B_4H_{10} is:
- (a) 4120 (b) 4220 (c) 4012 (d) 3203
106. Match list I (compounds) with list II (structures), and select the correct answer using the codes given below.
- | List-I | List-II |
|--------------------------|------------------------------|
| (A) XeO_4 | (i) square planar |
| (B) BrF_4^- | (ii) tetrahedral |
| (C) SeCl_4 | (iii) distorted tetrahedral. |
| (a) (A–ii) (B–iii) (C–i) | (b) (A–iii) (B–i) (C–ii) |
| (c) (A–ii) (B–i) (C–iii) | (d) (A–i) (B–ii) (C–iii) |
107. In the $\text{trans-PtCl}_2\text{L}(\text{CO})$ complex, the CO stretching frequency for $\text{L} = \text{NH}_3$, pyridine, NMe_3 decreases in the order.
- (a) pyridine > NH_3 > NMe_3 (b) NH_3 > pyridine > NMe_3
(c) NMe_3 > NH_3 > pyridine (d) pyridine > NMe_3 > NH_3
108. For the nuclear reactions.
- (A) ${}^8_4\text{Be} \longrightarrow 2{}^4_2\text{He}$ (B) ${}^{80}_{36}\text{Kr} \rightarrow 2{}^{40}_{18}\text{Ar}$
- (Given masses: ${}^8\text{Be} = 8.005300$, ${}^4\text{He} = 4.002603$ and ${}^{80}\text{Kr} = 79.81638$, ${}^{40}\text{Ar} = 39.96238$)
- The correct statement is:
- (a) (A) and (B) are both spontaneous fission processes.
(b) (A) is spontaneous fission but (B) is not.
(c) (B) is spontaneous fission but (A) is not
(d) Both (A) and (B) are not spontaneous fission processes.

109. A metal ion that replace manganese (II) ion in mangano-proteins without changing its function, is
 (a) Fe (II) (b) Zn (II) (c) Mg (II) (d) Cu (II)
110. In $^{57}\text{Fe}^*$ Mossbauer experiment, source of 14.4 keV (equivalent to 3.48×10^{12} MHz) is moved towards absorber at a velocity of 2.2 mm s^{-1} . The shift in frequency of the source for this sample is:
 (a) 35.5 MHz (b) 25.5 MHz (c) 20.2 MHz (d) 15.5 MHz
111. Bayer's process involves.
 (a) Synthesis of B_2H_6 from NaBH_4 (b) Synthesis of NaBH_4 from borax
 (c) Synthesis of NaBH_4 from B_2H_6 (d) Synthesis of $\text{B}_3\text{N}_3\text{H}_6$ from B_2H_6 .
112. A true statement about base hydrolysis of $[\text{Co}(\text{NH}_3)_5\text{Cl}]^{2+}$ is:
 (a) It is a first order reaction
 (b) The rate determining step involves the dissociation of chloride in $[\text{Co}(\text{NH}_3)_4(\text{NH}_2)\text{Cl}]^+$.
 (c) The rate is independent of the concentration of the base
 (d) The rate determining step involves the abstraction of a proton from $[\text{Co}(\text{NH}_3)_5\text{Cl}]^{2+}$.
113. The catalyst involved in carrying out the metathesis of 1-butene to give ethylene and 3-hexene is:
 (a)  (b) Na_2PdCl_4 (c) $\text{Co}_2(\text{CO})_8, \text{H}_2$ (d) $\text{RhCl}(\text{PPh}_3)_3$
114. The correct order of d-orbital splitting in a trigonal bipyramidal geometry is:
 (a) $d_{z^2} > d_{xz} > d_{x^2-y^2}, d_{xy}$ (b) $d_{xz}, d_{yz} > d_{x^2-y^2}, d_{xy} > d_{z^2}$
 (c) $d_{x^2-y^2}, d_{xy} > d_{z^2} > d_{xz}, d_{yz}$ (d) $d_{z^2} > d_{x^2-y^2} > d_{xy} > d_{xz}, d_{yz}$
115. For the following outer sphere electron transfer reactions.

$$[\text{Co}(\text{NH}_3)_6]^{2+} + [\text{Co}^*(\text{NH}_3)_6]^{3+} \rightarrow [\text{Co}(\text{NH}_3)_6]^{3+} + [\text{Co}^*(\text{NH}_3)_6]^{2+}$$

$$[\text{Ru}(\text{NH}_3)_6]^{2+} + [\text{Ru}^*(\text{NH}_3)_6]^{3+} \rightarrow [\text{Ru}(\text{NH}_3)_6]^{3+} + [\text{Ru}^*(\text{NH}_3)_6]^{2+}$$
 the rate constants are $10^{-6} \text{ M}^{-1} \text{ s}^{-1}$ and $8.2 \times 10^2 \text{ M}^{-1} \text{ s}^{-1}$ respectively. This difference in the rate constants is due to
 (a) A change from high spin to low spin in Co^* and high spin to low spin in Ru.
 (b) A change from high spin to low spin in Co^* and low spin to high spin Ru*.
 (c) A change from low spin to high spin in Co^* and the low spin state remains unchanged in Ru.
 (d) A change from low spin to high spin in Co^* and high spin to low spin in Ru*.
116. The greater stability of $((\text{CH}_3)_3\text{C}-\text{CH}_2-)_4\text{Ti}(\text{A})$ compared to that of $((\text{CH}_3)_2\text{CH}-\text{CH}_2-)_4\text{Ti}(\text{B})$ is due to
 (a) Hyperconjugation present in complex (A)
 (b) β -hydride elimination is not possible in complex (A)
 (c) Steric protection of titanium from reactive species in complex (A)
 (d) The stronger nature of Ti-C bond in complex (A).
117. The coordination number and geometry of cerium in $[\text{Ce}(\text{NO}_3)_6]^{2-}$ are respectively,
 (a) 6 and octahedron (b) 6 and trigonal prism
 (c) 8 and cubic (d) 12 and icosahedron

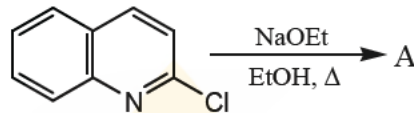
118. A compound **A** having the composition $\text{FeC}_9\text{H}_8\text{O}_3$ shows one signal at 2.5 ppm and another one around 5.0 ppm in its ^1H NMR spectrum. The IR spectrum of this compound shows two bands around and 1680 cm^{-1} . The compound follows the 18 electron rule of the following statements for **A**, the correct one is/are

- (A) It has $\eta^5\text{-Cp}$ group. (B) It has a terminal CO ligand.
 (C) It has a CH_3 ligand (D) It has Fe–H bond.
 (a) (A) and (B) only (b) (C) only (c) (A) and (C) only (d) (B) and (D) only.

119. In bacterial rubredoxin, the number of iron atoms, sulfur bridges and cysteine ligands are

	Fe atom	sulfur bridge	cysteine
(a)	4	4	4
(b)	2	2	4
(c)	2	2	2
(d)	1	0	4

120. In the following reaction, the product formed and the mechanism involved are



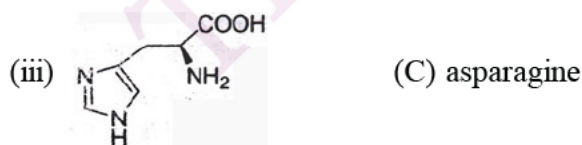
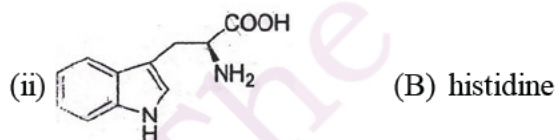
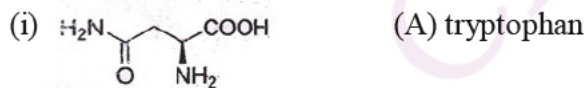
- (a) **A** is C1=CC=C2C=CC(=C1)N=C(OCC)C2 and is formed by addition-elimination mechanism.
 (b) **A** is C1=CC=C2C=CC(=C1)N=C(OCC)C2 and is formed by benzyne mechanism.
 (c) **A** is C1=CC=C2C=CC(=C1)N=C(OCC)C2 and is formed by benzyne mechanism.
 (d) **A** is C1=CC=C2C=CC(=C1)N=C(OCC)C2 and is formed by $\text{S}_{\text{N}}2$ displacement.

121. An optically active compound enriched with R-enantiomer (60% ee) exhibited $\alpha_{\text{D}} +$. If the

$[\alpha]_{\text{D}}$ value of the sample is -135° , the ratio of R and S enantiomers would be

- (a) R:S = 1:19 (b) R:S = 19:1 (c) R:S = 1:9 (d) R:S = 9:1

122. Match the amino acids with their structures:

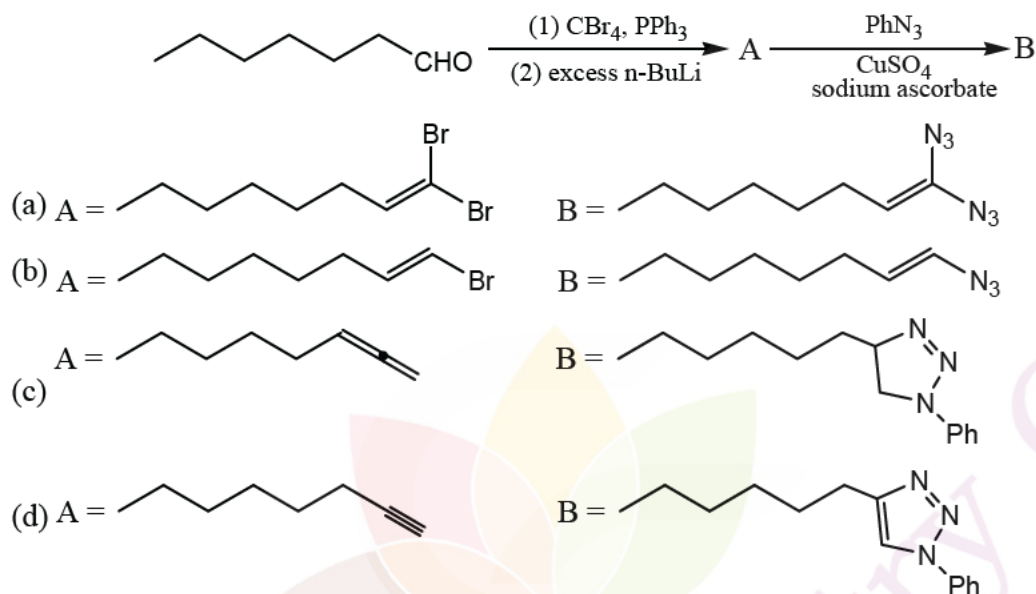


- (D) serine
(E) glutamic acid.

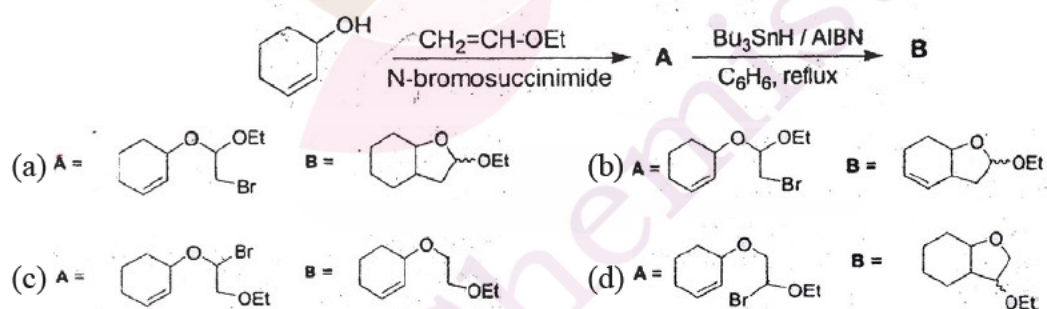
- (a) (i)-A (ii)-E (iii)-(C) (b) (i)-(C) (ii)-(D) (iii)-(B)
 (c) (i)-(A) (ii)-(B) (iii)-(D) (d) (i)-(C) (ii)-(A) (iii)-(B)

123. **Statement I.** U(VI) is more stable than Nd(VI).
Statement II. The valence electrons in U are in 5f, 6d and 7s orbitals.
 (a) Statements I and II are correct and Statement II is correct explanation of I.
 (b) Statements I and II are correct but Statement II is not an explanation for Statement I.
 (c) Statements I is correct and Statement II is incorrect.
 (d) Statements I and II both are incorrect.

124. The major products A and B in the following reaction sequence are



125. The major products A and B in the following reaction sequence are



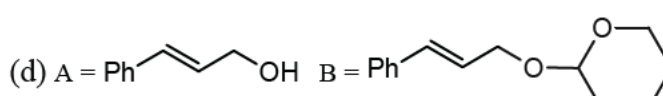
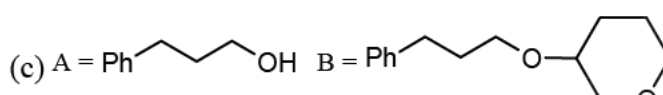
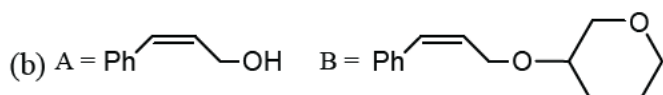
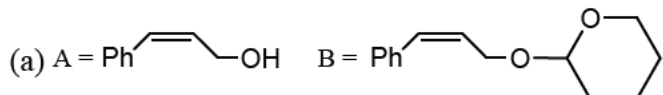
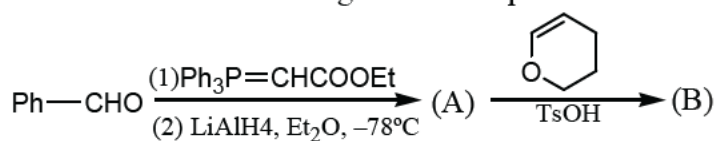
126. An organic compound having molecular formula $\text{C}_{15}\text{H}_{14}\text{O}$ exhibited the following ^1H and ^{13}C NMR spectral data.

^1H NMR : δ 2.4(s), 7.2(d, $J = 8$ Hz), 7.7(d, $J = 8$ Hz)

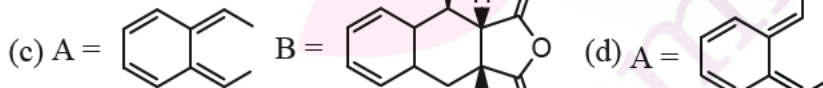
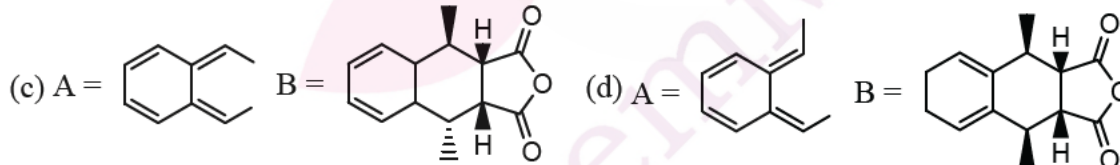
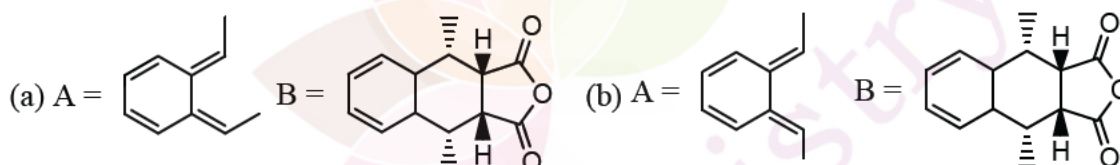
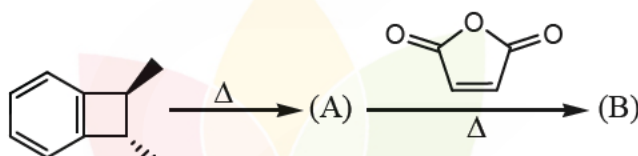
^{13}C NMR : δ 21.0, 129.0, 130.0, 136.0, 141.0, 190.0



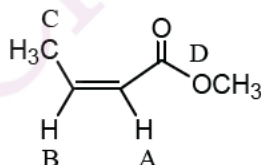
131. The major products A and B in the following reaction sequence are



132. The major products A and B in the following reaction sequence are



133. Appropriate ¹H NMR chemical shifts (δ) for the protons A-D for the following compound are



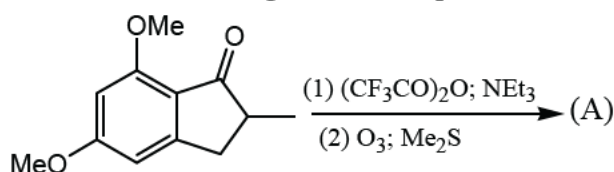
(a) A- 6.8; B-5.7; C-3.9; D -2.1ppm

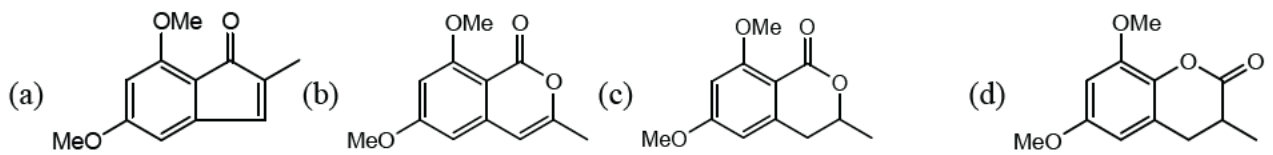
(b) A- 6.8; B-5.7; C-2.1; D -3.9 ppm

(c) A-5.7; B-6.8; C-3.9; D -2.1 ppm

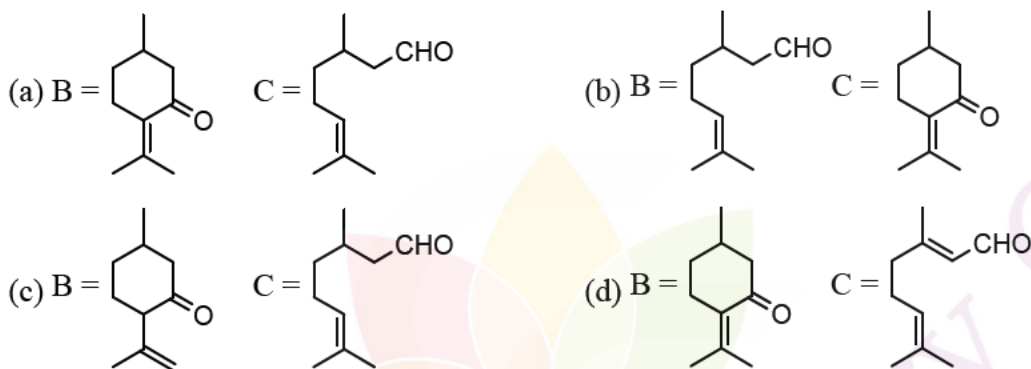
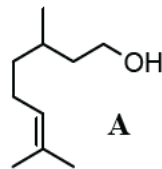
(d) A- 5.7; B - 6.8; C-2.1; D -3.9 ppm

134. The major product formed in the following reaction sequence is:

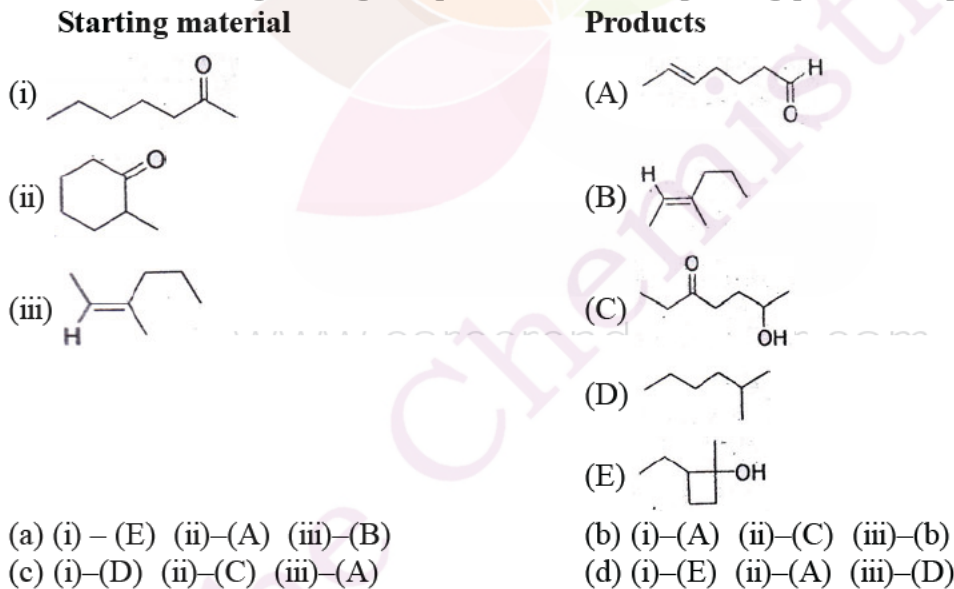




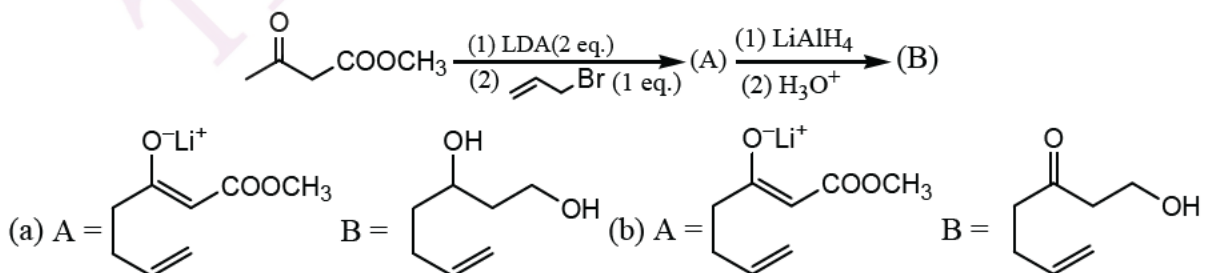
135. Citronellol A on oxidation with pyridinium chlorochromate (PCC) followed by treatment with aq. sodium hydroxide gives the product B (IR : 1680 cm^{-1}); whereas oxidation with PCC in the presence of sodium acetate gives product C (IR: 1720 cm^{-1}). Compound B and C are

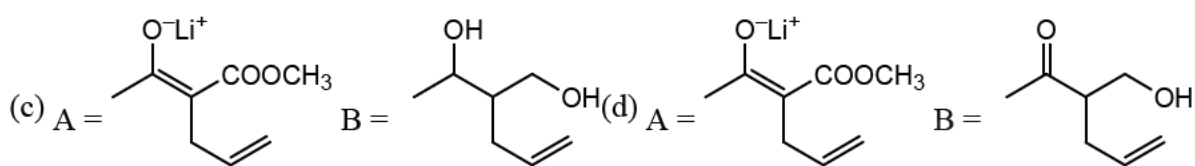


136. Match the following starting compounds with corresponding products in photochemical reactions:

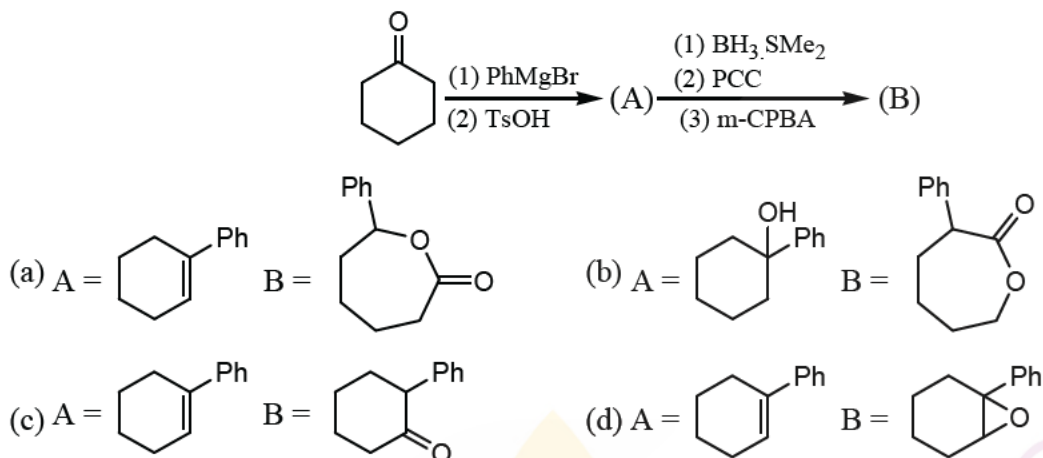


137. The major products A and B in the following reaction sequence are

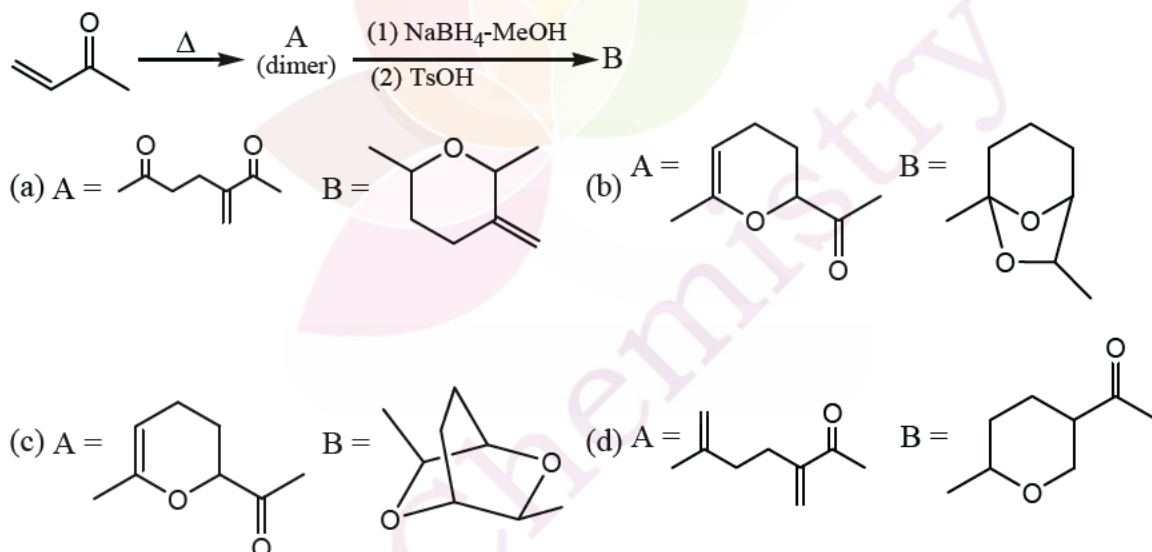




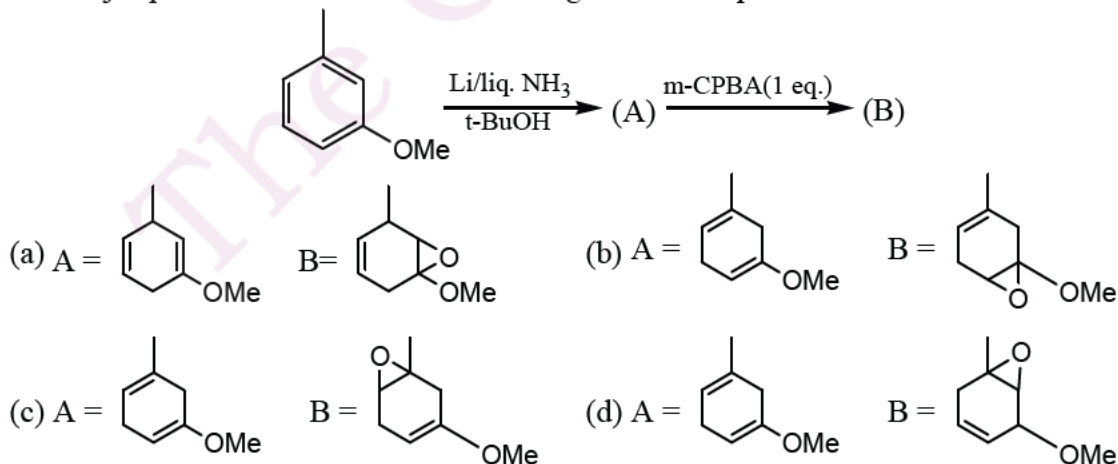
138. The major products A and B of the following reaction sequence are



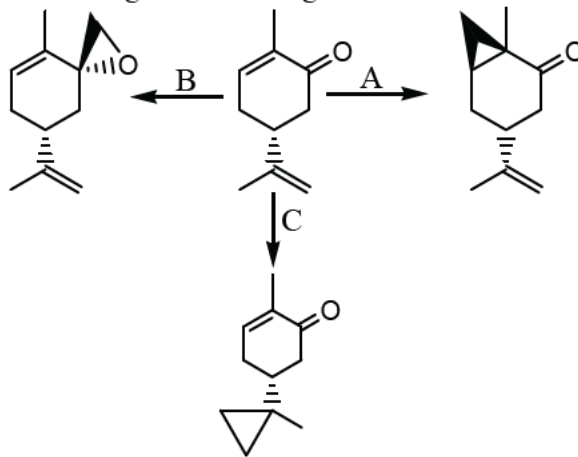
139. The major products A and B in the following reaction sequence are



140. The major products A and B in the following reaction sequence are

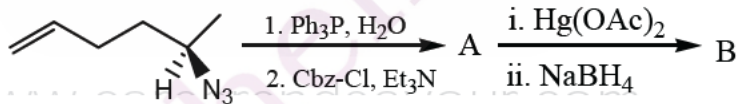


141. The correct reagents for effecting the following reactions are



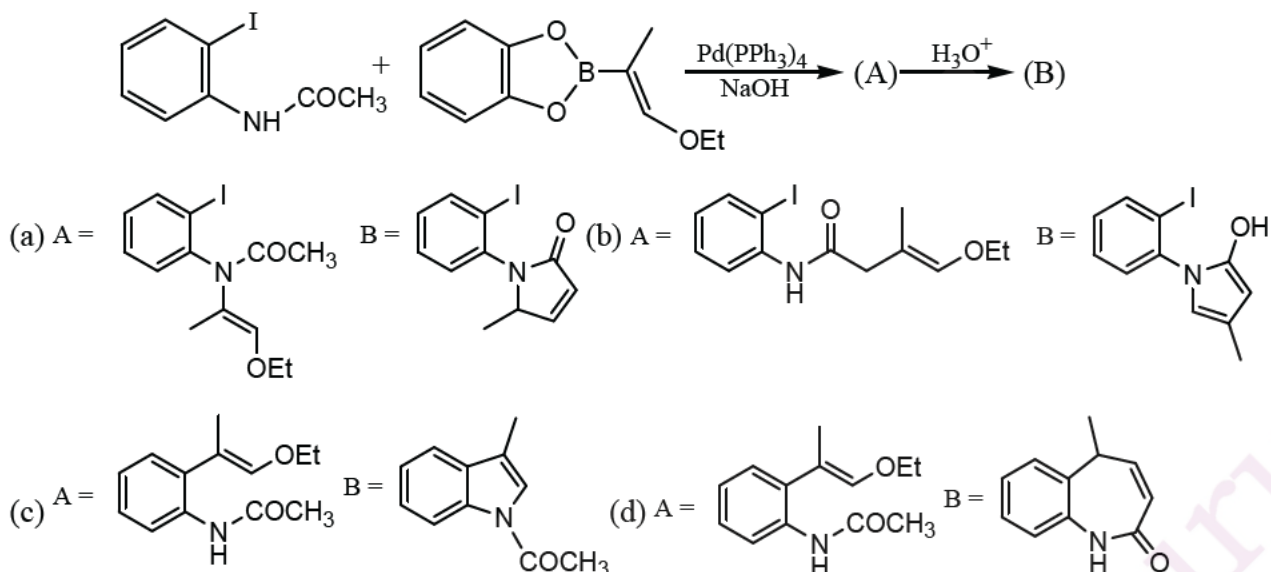
- (a) $A = \text{Me}-\overset{\oplus}{\text{S}}(\text{Me})_2-\text{CH}_2^-$ $B = \text{Me}-\overset{\oplus}{\text{S}}(\text{Me})_2-\text{CH}_2^-$ $C = \text{CH}_2\text{I}_2, \text{Zn-Cu}$
- (b) $A = \text{Me}-\overset{\ominus}{\text{S}}(\text{Me})_2-\text{CH}_2^+$ $B = \text{Me}-\overset{\oplus}{\text{S}}(\text{Me})_2-\text{CH}_2^-$ $C = \text{CH}_2\text{I}_2, \text{Zn-Cu}$
- (c) $A = \text{CH}_2\text{I}_2, \text{Zn-Cu}$ $B = \text{Me}-\overset{\oplus}{\text{S}}(\text{Me})_2-\text{CH}_2^-$ $C = \text{Me}-\overset{\oplus}{\text{S}}(\text{Me})_2-\text{CH}_2^-$
- (d) $A = \text{Me}-\overset{\oplus}{\text{S}}(\text{Me})_2-\text{CH}_2^-$ $B = \text{Me}-\overset{\oplus}{\text{S}}(\text{Me})_2-\text{CH}_2^-$ $C = \text{Tebbe's reagent}$

142. The major product A and B of the following reaction sequence are

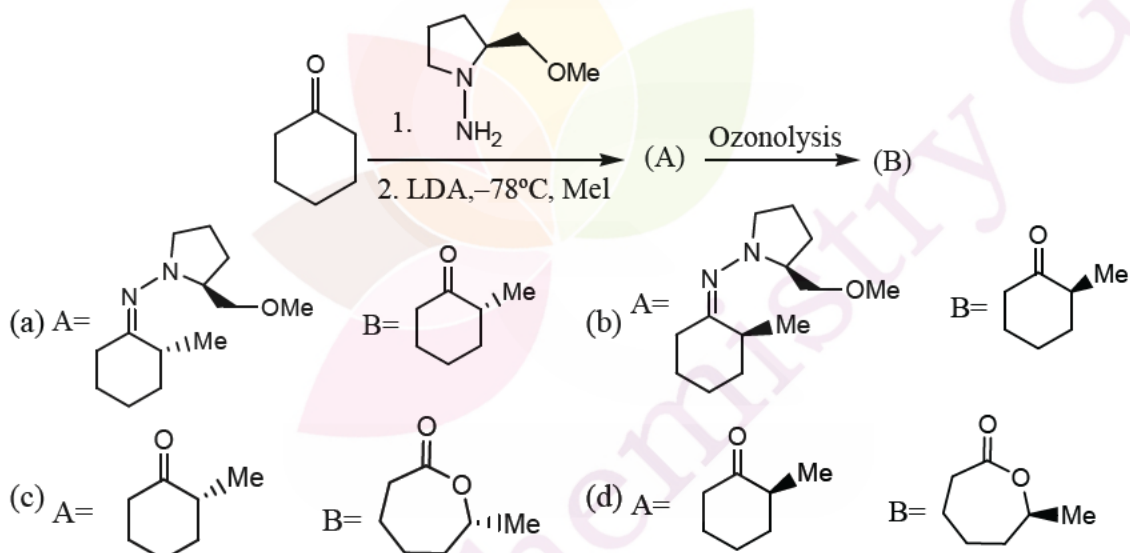


- (a) $A = \text{CH}_2=\text{CH}-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{NH}_2$ $B = \text{Cyclohexane ring with NH and methyl groups}$
- (b) $A = \text{CH}_2=\text{CH}-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{NHCOOBn}$ $B = \text{Cyclohexane ring with NHCOOBn and methyl groups}$
- (c) $A = \text{CH}_2=\text{CH}-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{NHCOOt-Bu}$ $B = \text{Cyclohexane ring with NHCOOt-Bu and methyl groups}$
- (d) $A = \text{CH}_2=\text{CH}-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{NHCOOt-Bu}$ $B = \text{Cyclohexane ring with NHCOOt-Bu and methyl groups}$

143. The major products A and B in the following synthetic sequence are



144. The major products A and B in the following synthetic strategy are



145. The product formed and the process involved in the following reaction are

