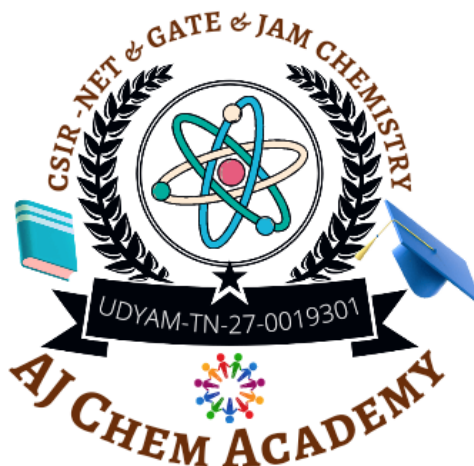


## GATE – 2024 – Chemistry



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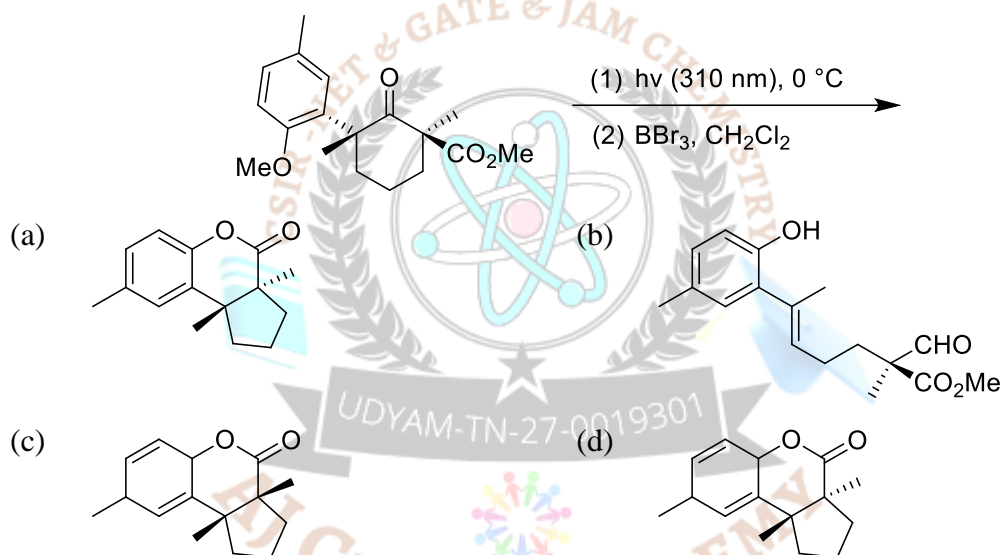
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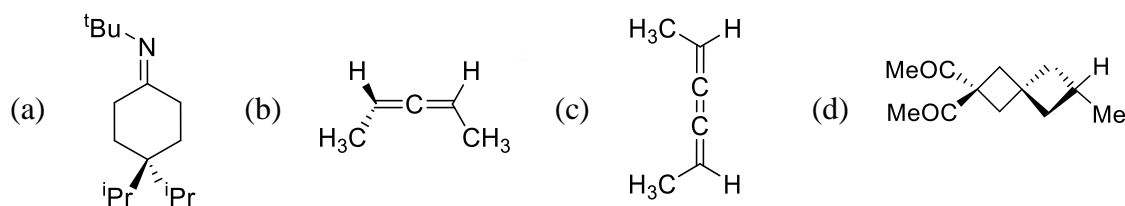


**(Q1-25) MCQ & NAT, carry ONE mark each (for each wrong answer: - 1/3). (\*\* No Negative Marks for NAT)**

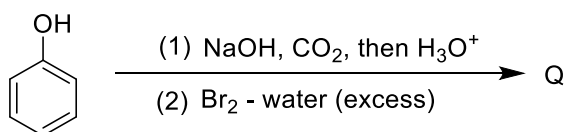
- Among the following, the compound with the **highest CO stretching frequency** is  
 (a)  $[\text{Mn}(\text{CO})_6]^+$  (b)  $[\text{V}(\text{CO})_6]^-$  (c)  $[\text{Cr}(\text{CO})_5]$  (d)  $[\text{Cr}(\text{dien})(\text{CO})_3]$
- The **ground state of  $[\text{Cr}(\text{H}_2\text{O})_6]^{2+}$**  is  
 (a)  $^5\text{E}_g$  (b)  $^5\text{T}_{2g}$  (c)  $^6\text{A}_{1g}$  (d)  $^6\text{A}_{2g}$
- The reaction of  **$\text{XeF}_2$  with  $\text{HN}(\text{SO}_2\text{F})_2$  at 273 K in  $\text{CF}_2\text{Cl}_2$  solvent yields**  
 (a)  $\text{XeF}_4 + \text{SO}_2 + \text{NH}_3$  (b)  $\text{Xe} + \text{SO}_2 + \text{N}_2 + \text{HF}$   
 (c)  $\text{SOF}_2 + \text{XeO}_2 + \text{NH}_3$  (d)  $\text{FXeN}(\text{SO}_2\text{F})_2 + \text{HF}$
- The **major product** in the following reaction sequence is



- Among the following, the **chiral compound** is



- The **major product** in the given reaction sequence is **Q**. The **mass spectrum of Q** shows **([M] = molecular ion peak)**

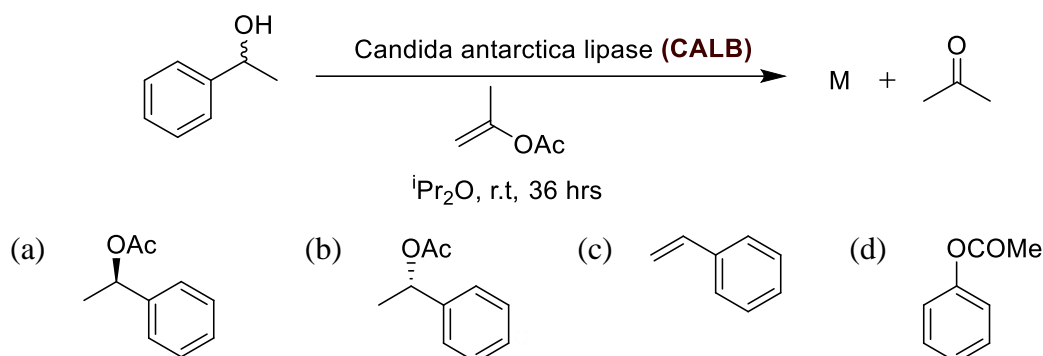


- (a)  $[\text{M}]$ ,  $[\text{M} + 2]$ ,  $[\text{M} + 4]$ , and  $[\text{M} + 6]$  peaks with relative intensity of 1: 1: 1: 1  
 (b)  $[\text{M}]$ ,  $[\text{M} + 2]$ ,  $[\text{M} + 4]$ , and  $[\text{M} + 6]$  peaks with relative intensity of 1: 3: 3: 1

(c) [M], [M + 2], and [M + 4] peaks with relative intensity of 1: 2: 1

(d) M and [M + 2] peaks with relative intensity of 1: 1

7. The **product M** in the following reaction is



8. **Critical micellar concentration** of a surfactant is **0.008 M** in water at **25°C**. If the **aggregation number** of the micelles is **80**, the concentration of the micelles (**in M**) present in **0.088 M** aqueous solution of the surfactant at **25°C** is

(a) 0.010 (b) 0.001 (c) 0.008 (d) 0.088

9. The **order** and the **number of classes** present in a group with the **irreducible representations** **A<sub>1</sub>, A<sub>2</sub>, B<sub>1</sub>, B<sub>2</sub>, E<sub>1</sub> and E<sub>2</sub>**, are, respectively,

(a) 6 and 6 (b) 12 and 6 (c) 6 and 3 (d) 12 and 3

10. The molecule **XY<sub>2</sub>** is microwave active and its **vibration-rotation spectrum** shows only **P** and **R** transitions. In the correct structure,

(a) X is the central atom in linear XY<sub>2</sub> (b) X is the central atom in bent XY<sub>2</sub>  
 (c) Y is the central atom in linear XY<sub>2</sub> (d) Y is the central atom in bent XY<sub>2</sub>

11. The complex(es) with **distorted octahedral structure** is (are)

(a) [VF<sub>6</sub>]<sup>3-</sup> (b) [FeF<sub>6</sub>]<sup>3-</sup> (c) [MnF<sub>6</sub>]<sup>3-</sup> (d) [Fe(CN)<sub>6</sub>]<sup>4-</sup>

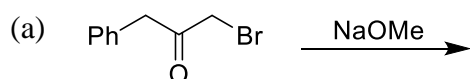
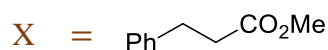
12. The compound(s) which show(s) the **perovskite structure** in solid state is (are)

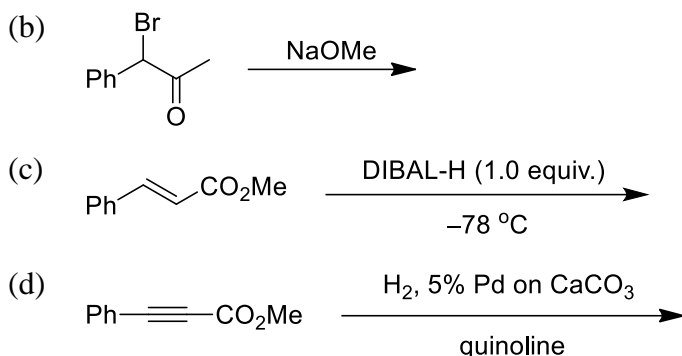
(a) CaTiO<sub>3</sub> (b) NiFe<sub>2</sub>O<sub>4</sub> (c) Fe<sub>3</sub>O<sub>4</sub> (d) CsPbI<sub>3</sub>

13. Among the following **metalloproteins**, the pair(s) of **non-heme proteins** is (are)

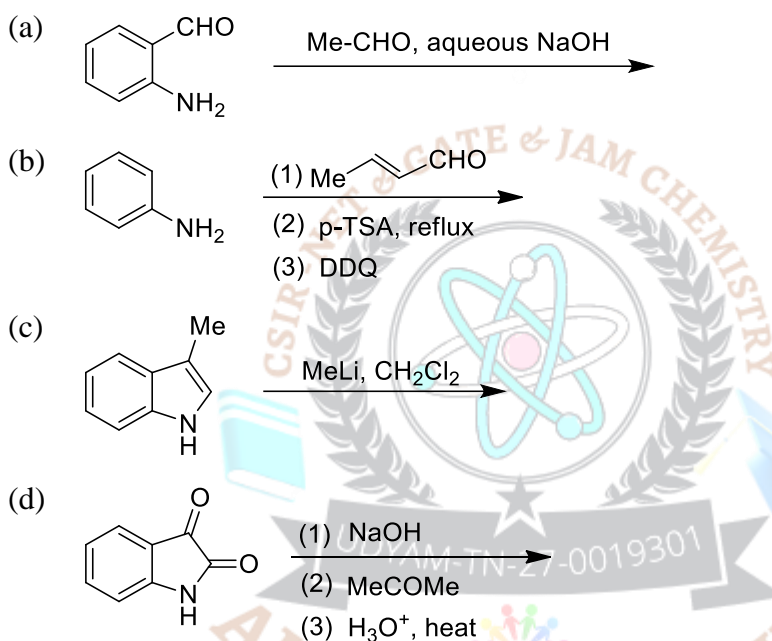
(a) Hemoglobin and Myoglobin (b) Hemocyanin and Carboxypeptidase  
 (c) Hemerythrin and Carbonic anhydrase (d) Cytochrome P-450 and hemocyanin

14. The reaction(s) that yield(s) **X** as the **major product** is (are)





15. The reaction(s) that yield(s) **2-methylquinoline** as the major product is (are)



16. The correct statement(s) for **decalin** is (are)

- (a) cis-Decalin is thermodynamically less stable than trans-decalin
- (b) cis-Decalin contains plane of symmetry
- (c) trans-Decalin undergoes ring inversion
- (d) trans-Decalin belongs to the point group of  $C_{2h}$

17. The correct statement(s) about  **$^4D_{5/2}$**  state of an atom is (are)

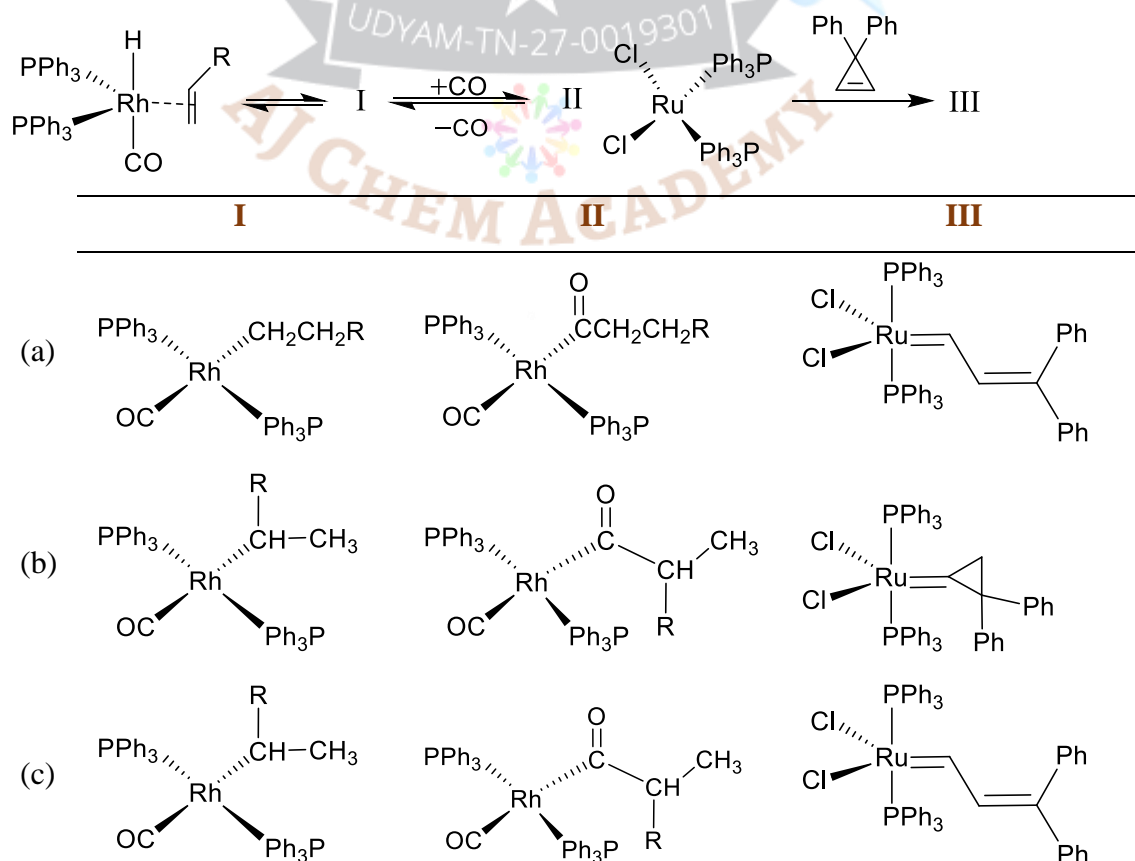
- (a) it corresponds to  $L = 2, S = 1/2$ , and  $J = 5/2$
- (b) it can originate from  $s^1p^2$  electronic configuration
- (c) it splits into five levels in the presence of magnetic field
- (d) it can show spectral transition to  $^4P_{3/2}$  state

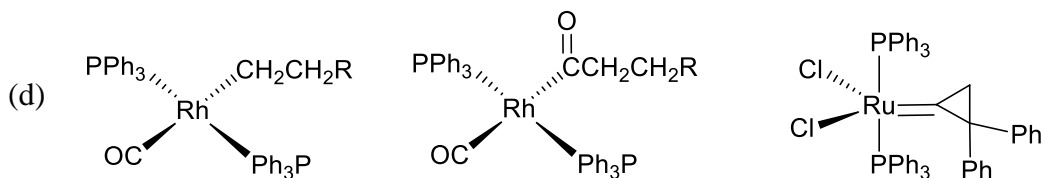
18. The correct statement(s) related to an **ensemble** is (are):

- (a) an ensemble is a collection of an infinite number of imaginary replications of the system of interest

- (b) all members of an ensemble are macroscopically identical and also have identical microstates
- (c) an ensemble average of any macroscopic property of the system is equal to the value of the property averaged over a sufficiently long time
- (d) all systems in a canonical ensemble need NOT have the same composition
19. The **non-dissociative adsorption** of a gas on a given surface at a fixed temperature follows **Langmuir isotherm**. The plot(s) which give(s) a **straight line** is (are)
- [Given:  $V$  = volume of the adsorbed gas,  $P$  = pressure of the gas]
- (a)  $1/V$  versus  $1/P$       (b)  $P/V$  and  $P$       (c)  $V$  versus  $P$       (d)  $V$  versus  $1/P$
20. The **crystal field stabilization energy** of  $[\text{Cr}(\text{NH}_3)_6]^{3+}$  with  $\Delta_o$  value of  $21600 \text{ cm}^{-1}$  is  $y \text{ cm}^{-1}$ . The value of  $|y|$  is \_\_\_\_\_. (rounded off to the nearest integer)
21. The **number of metal-metal bond(s)** in the complex  $[(^5\eta\text{-Cp})\text{Mo}(\text{CO})_2]_2$  is  $x$  and in  $[(^5\eta\text{-Cp})_2\text{Fe}_2(\text{CO})_3]$  is  $y$ . The value of  $x + y$  is \_\_\_\_\_. (Assume 18 electron rule is followed.)
22.  $^1\text{H-NMR}$  spectrum of a mixture containing  $\text{CH}_3\text{Br}$  ( $x$  mol) and  $(\text{CH}_3)_3\text{CBr}$  ( $y$  mol) shows **two singlets** at **2.7 ppm** and **1.8 ppm**, with the relative ratio of **3:1** (integration value), respectively. The value of  $x/y$  is \_\_\_\_\_. (rounded off to the nearest integer)
23. The value of  $\frac{e^2}{2\pi\epsilon_0 a_0}$  in **atomic unit of energy** is \_\_\_\_\_. (rounded off to the nearest integer) (e: charge of electron;  $a_0$ : Bohr radius;  $\epsilon_0$ : permittivity of vacuum)
24. The **partial vapor pressure** of **0.1 molal** solution of **B** in liquid **A** is **60 kPa** at **300 K**. The **partial vapor pressure** (in kPa) of a solution containing **B** with **mole fraction** of **0.1** in liquid **A** at **300 K** is \_\_\_\_\_. (rounded off to three decimal places) (Assume the solute **B** obeys Henry's law. The molar mass of **A** is  $80 \text{ g mol}^{-1}$ .)
25. Consider the following **two parallel irreversible first-order reactions**,
- $$\text{P1} \xleftarrow{k_1} \text{R} \xrightarrow{k_2} \text{P2} \quad (k_1 \text{ and } k_2 \text{ are the rate constants})$$
- where  $k_1 = 2k_2$  at **300 K**. After complete conversion of **R** at **300 K**, the concentration of **P1** in the reaction mixture was  $15 \text{ mol L}^{-1}$ . The initial concentration of **R** (in  $\text{mol L}^{-1}$ ) was \_\_\_\_\_. (rounded off to one decimal place)
- (Q26-55) MCQ & NAT, carry TWO marks each (for each wrong answer: - 2/3). (\*\* No Negative Marks for NAT)

26. Borax on treatment with NaOH and H<sub>2</sub>O<sub>2</sub> forms X. The compound X on reaction with PhCN at 60°C in methanol-water mixture gives Y as the major product. X and Y, respectively, are
- (a) NaB(O)(OH)<sub>2</sub>·nH<sub>2</sub>O and PhCONH<sub>2</sub>  
 (b) NaB(O)(OH)<sub>2</sub>·nH<sub>2</sub>O and PhCOOH  
 (c) Na<sub>2</sub>B<sub>2</sub>(O<sub>2</sub>)<sub>2</sub>(OH)<sub>4</sub>·nH<sub>2</sub>O and PhCONH<sub>2</sub>  
 (d) Na<sub>2</sub>B<sub>2</sub>(O<sub>2</sub>)<sub>2</sub>(OH)<sub>4</sub>·nH<sub>2</sub>O and PhCOOH
27. In the EPR spectrum of an aqueous solution of VOSO<sub>4</sub> at room temperature, the total number of hyperfine splitting signals is
- (a) 3 (b) 7 (c) 5 (d) 8
28. The hapticity of allyl and Cp and the ligation mode of NO in the thermodynamically stable complexes [(η<sup>x</sup>-allyl)Ru(CO)<sub>2</sub>(NO)] and [(η<sup>y</sup>-Cp)Ru(CO)<sub>2</sub>(NO)], respectively, are (The hapticity of allyl and Cp are denoted by η<sup>x</sup> and η<sup>y</sup>, respectively)
- (a) (η<sup>3</sup>, NO-bent) and (η<sup>5</sup>, NO-linear) (b) (η<sup>3</sup>, NO-linear) and (η<sup>5</sup>, NO-bent)  
 (c) (η<sup>1</sup>, NO-bent) and (η<sup>3</sup>, NO-bent) (d) (η<sup>1</sup>, NO-bent) and (η<sup>5</sup>, NO-linear)
29. In the following reactions, the structures of I, II and III, respectively, are





30. Consider the following data of a compound:

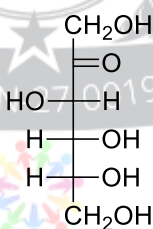
$^1\text{H-NMR}$  : 400 MHz,  $\text{DMSO-d}^6$

$\delta$  in ppm : 3.85(s, 6H), 6.73(t,  $J = 2.2$  Hz, 1H), 7.1(d,  $J = 2.2$  Hz, 2H) and 13.05(brs, 1H)

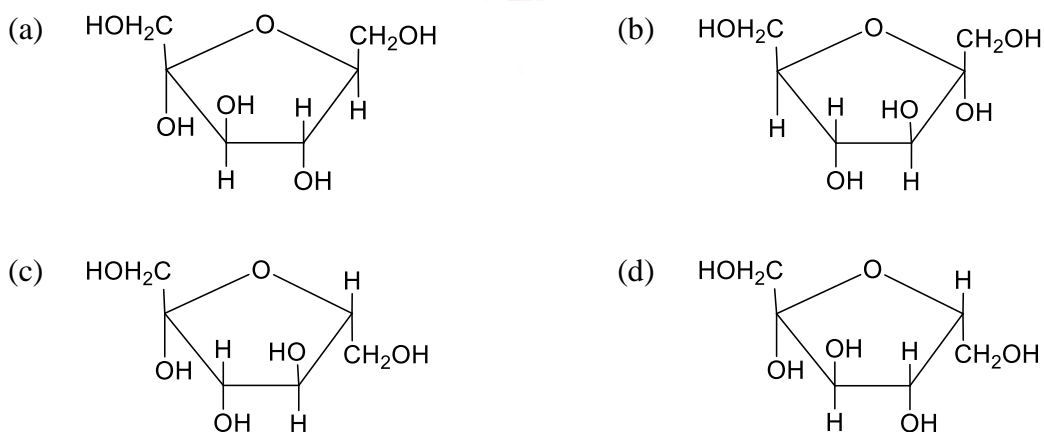
The compound is



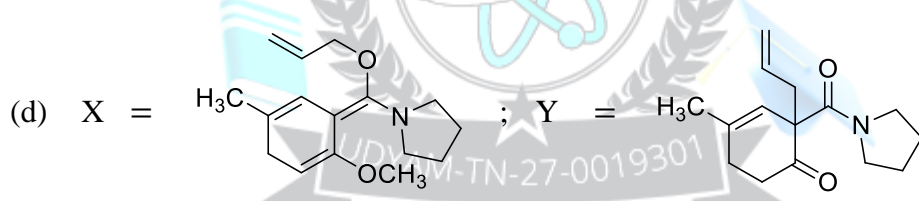
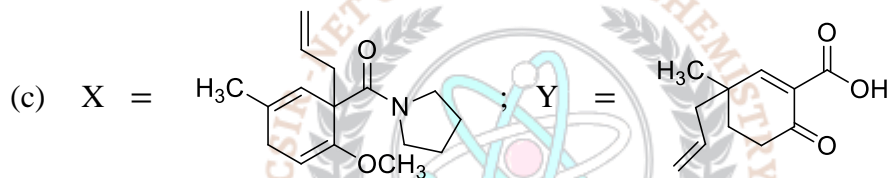
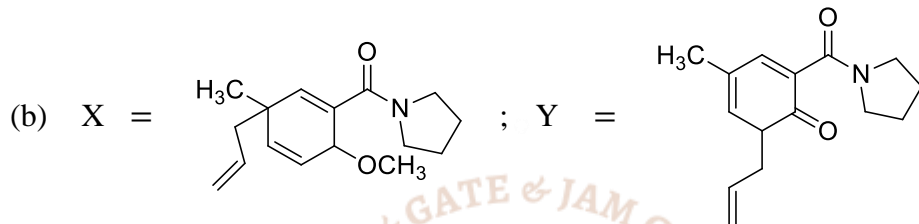
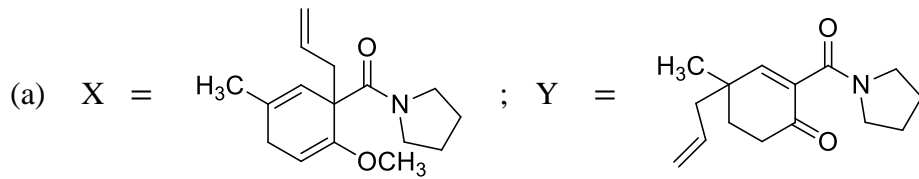
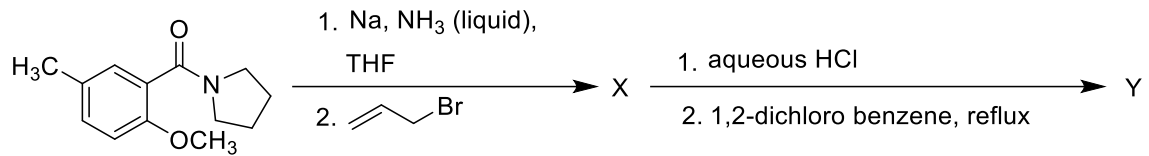
31. Fischer presentation of D-(–)-fructose is given below.



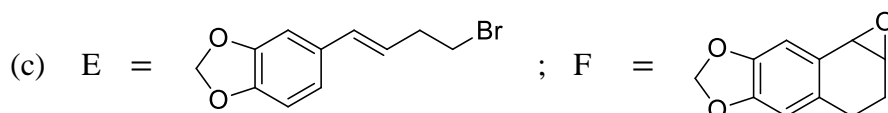
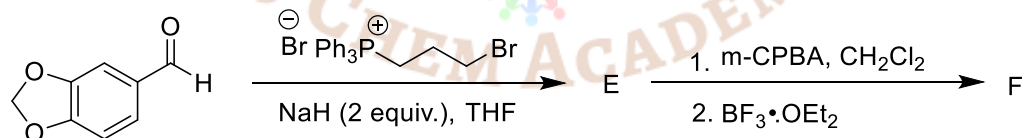
The correct structure of  $\alpha\text{-L-(+)-fructofuranose}$  is



32. The major products **X** and **Y** in the following reaction sequence are



33. The major products **E** and **F** in the following reaction sequence are



34.  $\psi_1, \psi_2, \psi_3$ , and  $\psi_4$  are four Hückel molecular orbitals of benzene with orbital energies  $E_1, E_2, E_3$  and  $E_4$ , respectively.

$$\psi_1 = \frac{1}{2}(\phi_B + \phi_C - \phi_E - \phi_F)$$

$$\psi_2 = 6^{-\frac{1}{2}}(\phi_A - \phi_B + \phi_C - \phi_D + \phi_E - \phi_F)$$

$$\psi_3 = 6^{-\frac{1}{2}}(\phi_A + \phi_B + \phi_C + \phi_D + \phi_E + \phi_F)$$

$$\psi_4 = 12^{-\frac{1}{2}}(2\phi_A + \phi_B - \phi_C - 2\phi_D - \phi_E + \phi_F)$$

The correct order of the orbital energies is (The six carbon atoms of benzene are denoted by A to F and  $\phi_j$  is the  $2p_z$  orbital of  $j^{\text{th}}$  carbon of benzene.)

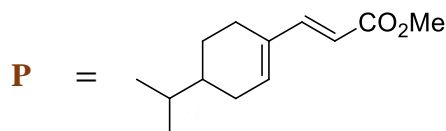
- (a)  $E_1 < E_2 = E_3 < E_4$  (b)  $E_4 < E_1 = E_3 < E_2$   
 (c)  $E_3 < E_1 = E_4 < E_2$  (d)  $E_3 < E_2 < E_1 = E_4$
35. Consider the following six vibrational modes:  
 Symmetric stretching of  $\text{CO}_2$ , O–H symmetric stretching of  $\text{H}_2\text{O}$ , stretching of  $\text{HCl}$ , stretching of  $\text{H}_2$ , N–H symmetric stretching of  $\text{NH}_3$ , and bending of  $\text{CO}_2$ . Among these modes, if  $k$  number of modes are IR active but Raman inactive,  $l$  number of modes are IR inactive but Raman active, and  $m$  number of modes are both IR and Raman active.  $k, l$  and  $m$ , respectively, are
- (a) 1, 3 and 2 (b) 3, 1 and 2 (c) 1, 2 and 3 (d) 2, 1 and 3
36. The correct statement for a thermally initiated radical polymerization in a solution is:

(Assume : Steady-state and equal reactivity of the propagating radicals, termination reactions are only by combination, and no chain transfer reaction.)

Given:  $R_p$  = rate of polymerization,  $DP$  = degree of polymerization,  $[I]$  = initiator concentration and  $[M]$  = monomer concentration)

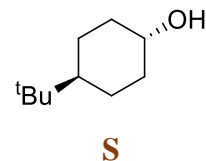
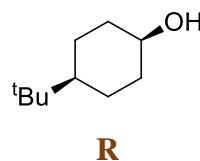
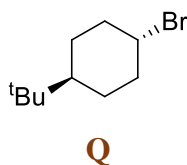
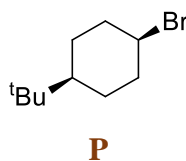
- (a) with increase in  $[I]$ , both  $R_p$  and  $DP$  increase  
 (b) with increase in  $[M]$ , both  $R_p$  and  $DP$  increase  
 (c)  $R_p$  decreases with increase in  $[I]$  but  $DP$  increases with increase in  $[M]$   
 (d)  $DP$  increases with increase in  $[I]$  and  $DP$  decreases with increase in  $[M]$ .
37. If  $q_t$  and  $Q_{t,m}$  are the molecular and molar translational partition functions of  $X_2$ , respectively, then  $\ln(Q_{t,m}) =$  (N is the Avogadro number)
- (a)  $N \ln q_t - N \ln N$  (b)  $N \ln q_t - \ln N$   
 (c)  $N \ln q_t + N \ln N + N$  (d)  $N \ln q_t - N \ln N + N$

38. Among the following, the **NMR active nucleus** (nuclei) is (are)  
 (a)  $^{12}\text{C}$  (b)  $^{19}\text{F}$  (c)  $^2\text{H}$  (d)  $^{16}\text{O}$
39. The complex(es) that exhibit(s) **optical isomerism** is (are)  
 (a)  $[\text{Fe}(\text{acac})_3]$  (b)  $\text{cis}-[\text{Co}(\text{en})_2\text{Cl}_2]^+$  (c)  $\text{trans}-[\text{Co}(\text{en})_2\text{Cl}_2]^+$  (d)  $[\text{Co}(\text{en})_3]^{3+}$
40. In aqueous solution of  $\text{K}_4[\text{Fe}(\text{CN})_6]$ , the allowed transition(s) is (are)  
 (a)  $^5\text{T}_{2g}$  to  $^3\text{E}_g$  (b)  $^1\text{A}_{1g}$  to  $^1\text{T}_{1g}$  (c)  $^1\text{A}_{1g}$  to  $^1\text{T}_{2g}$  (d)  $^5\text{T}_{2g}$  to  $^5\text{E}_g$
41. The correct option(s) that give(s) **P** as the **major product** is (are)



- (a)   
 (1) LDA, THF,  $-78^\circ\text{C}$ ,  $\text{Tf}_2\text{O}$   
 (2) Methyl acrylate,  $\text{Pd}(\text{Ph}_3\text{P})_2\text{Cl}_2$  (2 mol%),  $\text{Et}_3\text{N}$  (3 equiv.), DMF
- (b)   
 (1)  $\text{TsNHNH}_2$ , MeLi (2 equiv.), THF,  $-78^\circ\text{C}$   
 (2) DMF,  $0^\circ\text{C}$   
 (3) NaH,  $(\text{OMe})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{Me}$ , THF,  $0^\circ\text{C}$  to reflux
- (c)   
 (1)  $\text{Me}_3\text{SiCH}(\text{Li})\text{Cl}$ , THF,  $-78^\circ\text{C}$   
 (2)  $\text{HClO}_4$ , THF  
 (3) Zn,  $\text{BrCH}_2\text{CO}_2\text{Me}$   
 (4) p-TSA, reflux
- (d)   
 (1) L-Selectride, THF  
 (2) MsCl,  $\text{Et}_3\text{N}$ ; then NaCN  
 (3) DIBAL-H (1equiv.), THF,  $-78^\circ\text{C}$   
 (4)  $\text{Ph}_3\text{P}=\text{CH}-\text{CO}_2\text{Me}$

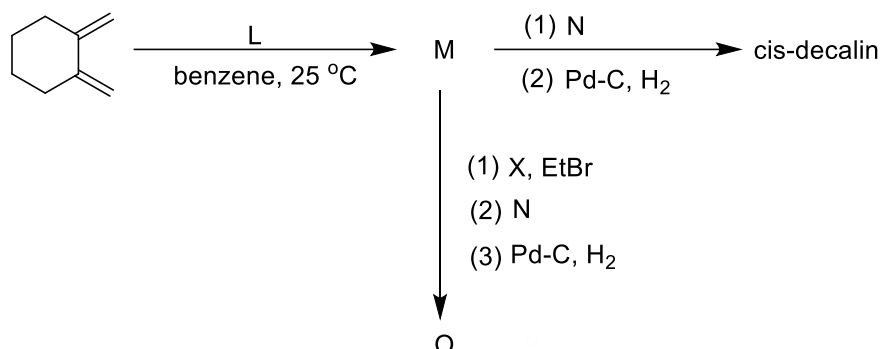
42. The correct statement(s) regarding **P, Q, R** and **S** is(are):



- (a) P reacts faster than Q with  $\text{PhSNa}$  in DMF as a solvent  
 (b) Q reacts faster than P with  $\text{NaN}_3$  in DMF as a solvent

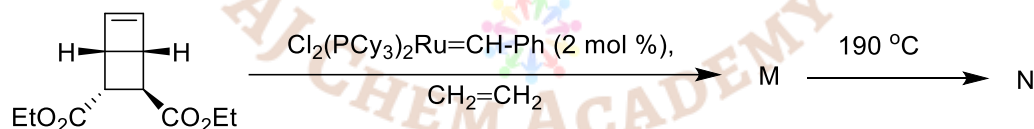
- (c) R reacts faster than S when treated with  $\text{TsCl}/\text{Et}_3\text{N}$  in DCM as a solvent  
 (d) R gets oxidized faster than S when treated with  $\text{CrO}_3$  in DCM as a solvent

43. Consider the following reaction sequence. The correct option(s) is (are)



- (a)  $\text{M} =$  ;  $\text{N} = \text{Na-Hg/MeOH}$   
 (b)  $\text{X} = \text{LDA}$  ;  $\text{O} =$    
 (c)  $\text{L} = \text{acrolein}$  ;  $\text{O} =$    
 (d)  $\text{X} = \text{LiAlH}_4$  ;  $\text{L} = (\text{Vinylsulfonyl})\text{benzene}$

44. Consider the following reaction sequence where **M** and **N** are the major products. The correct option(s) is (are)

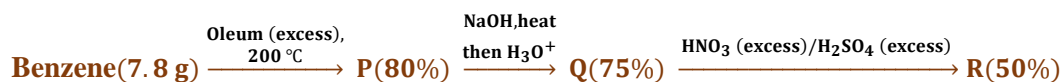


- (a)  $\text{M} =$    
 (b)  $\text{M} =$    
 (c)  $\text{N} =$    
 (d)  $\text{N} =$

45. The correct statement(s) about the relationship for the **H-atoms** in the following compounds is (are):



- (a)  $H_1$  and  $H_3$  are enantiotopic;  $H_2$  and  $H_3$  are diastereotopic  
 (b)  $H_1$  and  $H_3$  are diastereotopic;  $H_2$  and  $H_3$  are enantiotopic  
 (c)  $H_5$  and  $H_7$  are enantiotopic;  $H_6$  and  $H_7$  are homotopic  
 (d)  $H_5$  and  $H_7$  are homotopic;  $H_6$  and  $H_7$  are enantiotopic
46. Among the following, the **correct** statement(s) is (are):
- (a) The normalization factor of a Slater determinant for a 3-electron atom is  $\sqrt{\frac{1}{3}}$   
 (b) The number of nodes in the radial wave function of 3s orbital of a hydrogen atom is the same as the number of nodes in the angular wave function of a 4d orbital of hydrogen atom  
 (c) The energy separation between any two adjacent states is same for a harmonic oscillator, while it is different for a rigid rotor  
 (d) The magnitude of the total spin angular momentum of an  $\alpha$  electron is the negative of that of a  $\beta$  electron
47. Among the following, the **correct** statement(s) is (are):
- (a)  $C_2$  symmetry element is present in  $H_2O$  and  $H_2O_2$  but NOT in  $PCl_5$   
 (b) both  $C_2$  and  $C_3$  symmetry elements are present in  $CCl_4$  and  $SF_6$   
 (c) one  $\sigma_h$  and three  $\sigma_d$  symmetry elements are present in benzene  
 (d)  $\sigma_v$  symmetry element is present in  $NH_3$  but NOT in  $BF_3$
48.  $\Delta S^\circ$  (in  $J\ mol^{-1}\ K^{-1}$ ) for the given reaction at 298 K is \_\_\_\_\_. (rounded off to two decimal places)  
 $[Cu(H_2O)_6]^{2+} + en \rightleftharpoons [Cu(H_2O)_4(en)]^{2+} + 2H_2O$   
 (Given:  $\log K_1 = 10.6$ , Where  $K_1$  is the equilibrium constant.  $\Delta H^\circ = -54\ kJ\ mol^{-1}$  and  $R = 8.314\ J\ mol^{-1}\ K^{-1}$ )
49. The **turnover frequency** (in  $h^{-1}$ ) of a reaction where **5 mol% of a catalyst is required for 90% conversion in 3h** is \_\_\_\_\_. (rounded off to the nearest integer)
50. In **thermogravimetric analysis**, **12.45 mg of  $CuSO_4 \cdot 5H_2O$**  was subjected to heating under  $N_2$  atmosphere. At a particular temperature, there was a **weight loss of 3.6 mg**. The **number of water molecule(s) lost per formula unit** is \_\_\_\_\_. (rounded off to the nearest integer)  
 (Given: molar mass (in  $g\ mol^{-1}$ ) of  $H = 1.0$ ,  $O = 16.0$ ,  $S = 32.0$  and  $Cu = 63.5$ )
51. In the given reaction sequence, the **amount of 'R' produced (in g)** is \_\_\_\_\_.



(Given: molar mass (in  $\text{g mol}^{-1}$ ) of H = 1, C = 12, N = 14, O = 16 and S = 32)  
(rounded off to two decimal places)

52. The wave function of a particle in a cubic box (of side L) is given by

$$\psi(x, y, z) = \sqrt{32/L^3} \sin \frac{\pi x}{L} \cos \frac{\pi x}{L} \sin \frac{2\pi y}{L} \sin \frac{\pi z}{L}$$

The ratio of the energy of the state corresponding to the above wave function to the ground state energy is \_\_\_\_\_. (rounded off to the nearest integer)

53.  $\phi_1$  and  $\phi_2$  are normalized eigenfunctions of a Hermitian operator.

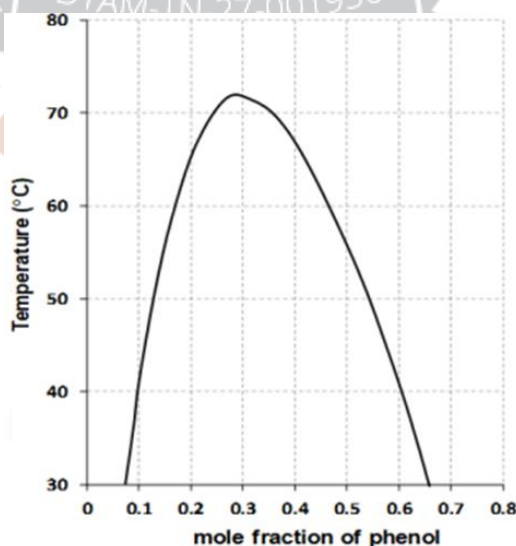
$$|\psi\rangle = 3i|\phi_1\rangle + 2|\phi_2\rangle \text{ and } |\chi\rangle = -2i|\phi_1\rangle + 5|\phi_2\rangle.$$

The value of  $\langle\psi|\chi\rangle + \langle\chi|\psi\rangle$  is \_\_\_\_\_. (rounded off to the nearest integer)

54. 2 mol of a monoatomic ideal gas with initial volume of 5 L and pressure 10 bar undergoes an irreversible adiabatic expansion against a constant final pressure of 1 bar. The final volume (in L) is \_\_\_\_\_. (rounded off to one decimal place)

(Given:  $R = 8.314 \times 10^{-2} \text{ L bar mol}^{-1} \text{ K}^{-1}$ )

55. The following figure shows an experimental liquid-liquid phase diagram of phenol and water at the vapor pressure of the system. The total amount of phenol and water (in mol) present in the phenol-rich phase when 5 mol of water was shaken with 5 mol of phenol at  $40^\circ\text{C}$  is \_\_\_\_\_. (rounded off to one decimal place)



### Answer Key

Q.No	Ans
1.	a
2.	a
3.	d

Q.No	Ans
21.	5
22.	9
23.	2

Q.No	Ans
41.	a & b
42.	a & d
43.	a & b

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4.	a
5.	b
6.	b
7.	a
8.	b
9.	b
10.	c
11.	a & c
12.	a & d
13.	b & c
14.	a & b
15.	b & d
16.	a & d
17.	d
18.	a & c
19.	a & b
20.	25920

24.	745 to 760
25.	22.5
26.	c
27.	d
28.	b
29.	a
30.	b
31.	a
32.	a
33.	a
34.	c
35.	c
36.	b
37.	d
38.	b & c
39.	a,b & d
40.	b & c

44.	a & c
45.	a & c
46.	b & c
47.	b & c
48.	21.40 to 22.00
49.	6
50.	4
51.	7.35
52.	3
53.	8 to 8
54.	31.8 to 32.2
55.	7.8 to 8.2

Q. 1 – 10	1 Mark (MCQ)
Q. 26 – 37	2 Mark (MCQ)

Q. 11 – 19	1 Mark (MSQ)
Q. 38 – 47	2 Mark (MSQ)

Q. 20-25	1 Mark (NAT)
Q. 48 – 55	2 Mark (NAT)

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