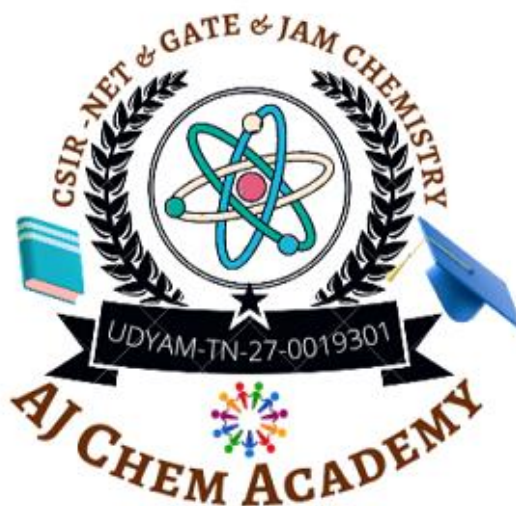


## GATE – 2019 – Chemistry



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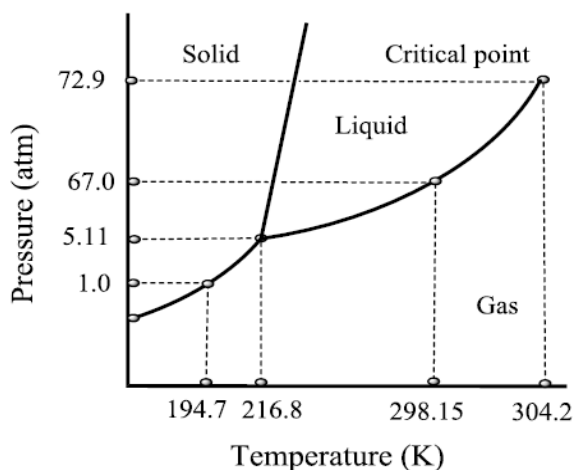


**Q.1 – Q.13 Multiple Choice Question (MCQ), carry ONE mark each (for each wrong answer: – 1/3).**

- The INCORRECT statement about the solid-state structure of CsCl and CaF<sub>2</sub> is:**
  - Cations in both solids exhibit coordination number 8
  - CsCl has bcc type structure and CaF<sub>2</sub> has cubic close pack structure
  - Radius ratio for Cs/Cl and Ca/F is 0.93 and 0.73, respectively
  - Both exhibit close pack structure
- The INCORRECT statement about the interhalogen compound ICl<sub>3</sub> is:**
  - It exists as a dimer
  - Geometry around the iodine is tetrahedral in solid-state
  - It decomposes as ICl and Cl<sub>2</sub> in gas-phase
  - Liquid ICl<sub>3</sub> conducts electricity
- Among the following carbon allotropes, the one with discrete molecular structure is**
  - Diamond
  - $\alpha$ -Graphite
  - $\beta$ -Graphite
  - Fullerene
- The INCORRECT statement about the silicones is:**
  - They are thermally unstable because of the Si–C bond
  - They are insoluble in water
  - They are organosilicon polymers
  - They have stable silica-like skeleton (–Si–O–Si–O–Si–)
- The  $\Delta_o$  value of  $[\text{Ni}(\text{H}_2\text{O})_6]^{2+}$  is  $8500 \text{ cm}^{-1}$ . The  $\Delta_o$  values for  $[\text{NiCl}_6]^{4-}$  and  $[\text{Ni}(\text{NH}_3)_6]^{2+}$  compared to  $[\text{Ni}(\text{H}_2\text{O})_6]^{2+}$  are**
  - higher and lower, respectively
  - lower and higher, respectively
  - higher in both complex ions
  - lower in both complex ions
- In Freundlich isotherm, a linear relationship is obtained in the plot of**  
**( $\theta$  = surface coverage and  $p$  = partial pressure of the gas)**
  - $\theta$  vs  $p$
  - $\ln(\theta)$  vs  $\ln(p)$
  - $\ln(\theta)$  vs  $p$
  - $\theta$  vs  $\ln(p)$
- Micelle formation is accompanied by the**
  - decrease in overall entropy due to ordering
  - increase in overall entropy mostly due to increase in solvent entropy
  - increase in overall entropy mostly due to increase in solute entropy
  - increase in overall entropy and decrease in enthalpy
- Consider the following phase diagram of CO<sub>2</sub> (not to scale). At equilibrium, the**

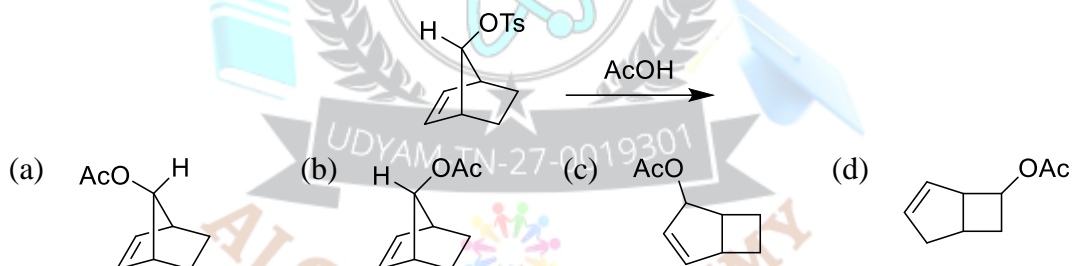


**INCORRECT** statement is:



- (a) At 200 K, on increasing the pressure from 1 to 50 atm, CO<sub>2</sub> gas condenses to liquid  
 (b) It is not possible to obtain liquid CO<sub>2</sub> from gaseous CO<sub>2</sub> below 5.11 atm  
 (c) Both liquid and gas phase of CO<sub>2</sub> coexist at 298.15 K and 67 atm  
 (d) With increasing pressure, the melting point of solid CO<sub>2</sub> increases

9. The **major product** formed in the following reaction is,

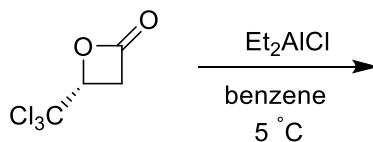


10. The **Woodward-Hoffmann condition** to bring out the following transformation is

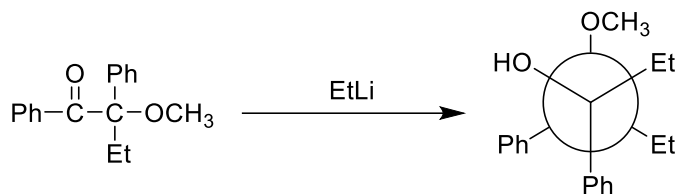


- (a)  $\Delta$ , conrotatory (b)  $\Delta$ , disrotatory (c)  $h\nu$ , disrotatory (d)  $h\nu$ , conrotatory

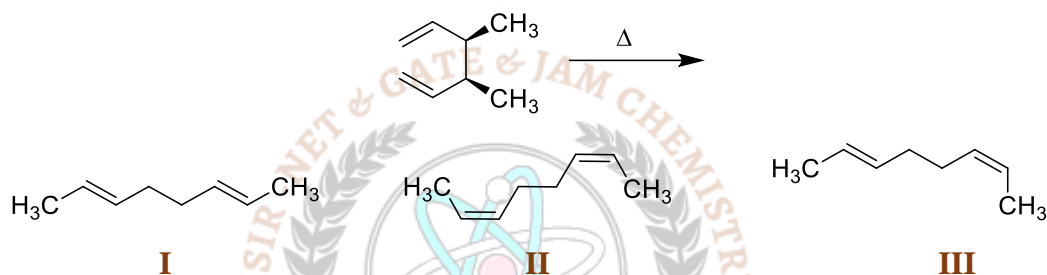
11. The **major product** formed in the following reaction is



12. In the following reaction, the **stereochemistry of the major product** is predicted by the



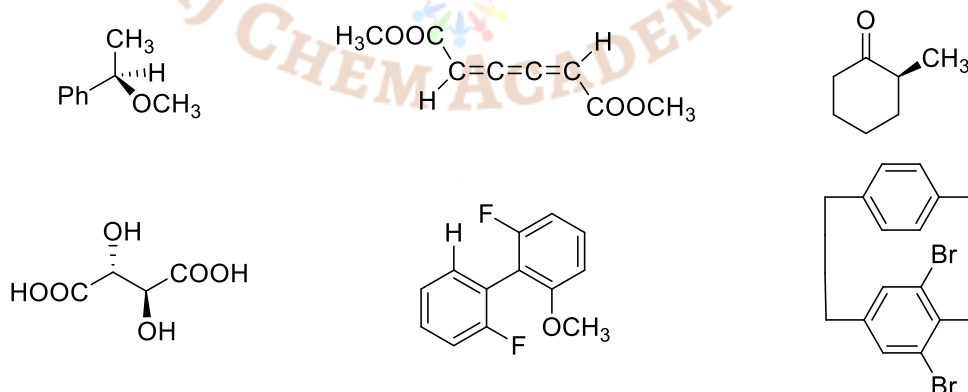
- (a) Cram's model  
(b) Cram's chelation model  
(c) Felkin model  
(d) Felkin-Anh model
13. The **product(s)** formed in the following reaction is (are)



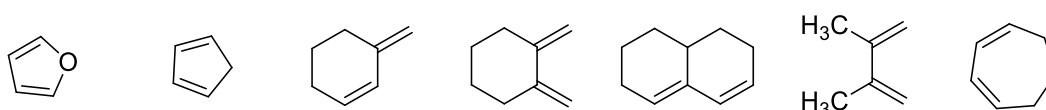
- (a) I only  
(b) II only  
(c) III only  
(d) mixture of I and II

**Q.14 – Q.25 Numerical Answer Type (NAT), carry ONE mark each (no negative marks).**

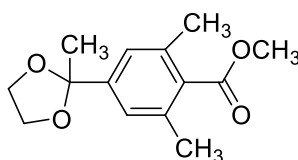
14. Among the following compounds, the number of compounds that **DO NOT** exhibit **optical activity at room temperature** is \_\_\_\_\_.



15. The number of following **diene(s)** that undergo **Diels-Alder reaction with methyl acrylate** is \_\_\_\_\_.



16. The number of  $^1\text{H-NMR}$  signals observed for the following compound is \_\_\_\_\_.



17. The number of CO stretching bands in IR spectrum of trigonal bipyramidal  $\text{Cis-M}(\text{CO})_3\text{L}_2$  is \_\_\_\_\_. ( $\text{M}$  = metal and  $\text{L}$  = monodentate ligand)
18. On heating a sample of 25 mg hydrated compound (molecular weight = 250 g/mol) in thermogravimetric analysis, 16 mg of dehydrated compound remains. The number of water molecules lost per molecule of hydrated compound is \_\_\_\_\_.  
(Molecular weight of water = 18 g/mol)
19. The total number of  $\alpha$  and  $\beta$  particles emitted in the given radioactive decay is \_\_\_\_.  

$${}_{92}^{238}\text{U} \rightarrow {}_{82}^{210}\text{Pb}$$
20. An ideal gas occupies an unknown volume  $V$  liters (L) at a pressure of 12 atm. The gas is expanded isothermally against a constant external pressure of 2 atm so that its final volume becomes 3 L. The work involved for this expansion process is \_\_\_\_\_ cal. (Round off to two decimal places)  
(Gas constant  $R = 0.082 \text{ L atm mol}^{-1}\text{K}^{-1} = 2 \text{ cal mol}^{-1}\text{K}^{-1}$ )
21. The entropy change for the melting of 'x' moles of ice (heat of fusion is  $80 \text{ cal g}^{-1}$ ) at 273 K and 1 atm pressure is  $28.80 \text{ cal K}^{-1}$ . The value of 'x' is \_\_\_\_\_. (Round off to two decimal places)  
(Molecular weight of water = 18 g/mol)
22. Consider a two-state system at thermal equilibrium having energies 0 and  $2k_{\text{B}}T$  for which the degeneracies are 1 and 2, respectively. The value of the partition function at the same absolute temperature  $T$  is \_\_\_\_\_. (Round off to two decimal places)  
( $k_{\text{B}}$  is the Boltzmann constant)
23. Consider a system of three identical and distinguishable non-interacting particles and three available nondegenerate single particle energy levels having energies 0,  $\epsilon$  and  $2\epsilon$ . The system is in contact with a heat bath of temperature  $T$  K. A total energy of  $2\epsilon$  is shared by these three particles. The number of ways the particles can be distributed is \_\_\_\_\_.
24. In a 400 MHz  ${}^1\text{H}$ -NMR spectrometer, a proton resonates at 1560 Hz higher than that of tetramethylsilane. The chemical shift value of this proton is \_\_\_\_\_ ppm.  
(Round off to one decimal place & Chemical shift of TMS is fixed at zero ppm)
25. Gas phase bond length and dipole moment of a compound (MX) is  $3 \text{ \AA}$  and  $10.8 \text{ D}$ , respectively. The ionic character in gas phase MX is \_\_\_\_\_ %.



(Round off to one decimal place &  $1D = 3.336 \times 10^{-30} \text{ C m}$ )

**Q.26 – Q.44 Multiple Choice Question (MCQ), carry TWO marks each (for each wrong answer: – 2/3).**

26. The experimentally observed magnetic moment values, which match well with the spin-only values for the pair of aqueous ions is  
(Atomic number: Cr = 24, Co = 27, Gd = 64, Tb = 65, Dy = 66 and Lu = 71)  
(a) Cr(III) and Gd(III) (b) Co(II) and Gd(III)  
(c) Cr(III) and Dy(III) (d) Lu(III) and Tb(III)
27. Among the following compounds, a normal spinel is  
(a)  $\text{MgFe}_2\text{O}_4$  (b)  $\text{ZnFe}_2\text{O}_4$  (c)  $\text{CoFe}_2\text{O}_4$  (d)  $\text{CuFe}_2\text{O}_4$
28. Following are the examples of silicate minerals  
Zircon,  $\text{ZrSiO}_4$       Beryl,  $\text{Be}_3\text{Al}_2\text{Si}_6\text{O}_{18}$       Pyrophyllite,  $\text{Al}_2(\text{OH})_2[(\text{Si}_2\text{O}_5)_2]$   
I                                      II                                      III  
The correct structural description of the minerals is  
I                                      II                                      III
- |     |                 |                   |                     |
|-----|-----------------|-------------------|---------------------|
| (a) | Ortho silicate  | ; Cyclic silicate | and Sheet silicate  |
| (b) | Ortho silicate  | ; Sheet silicate  | and Cyclic silicate |
| (c) | Cyclic silicate | ; Sheet silicate  | and Ortho silicate  |
| (d) | Sheet silicate  | ; Ortho silicate  | and Cyclic silicate |
29. In the EPR spectrum of a methyl radical, the number of lines and their relative intensities, respectively, are  
(a) 1 and 1 (b) 3 and 1:2:1 (c) 4 and 1:2:2:1 (d) 4 and 1:3:3:1
30. The product obtained in the reaction of  $\text{Mn}_2(\text{CO})_{10}$  with  $\text{Br}_2$  is  
(a)  $\text{Mn}(\text{CO})_5\text{Br}$  (b)  $\text{Mn}_2(\text{CO})_8\text{Br}_2$  (c)  $\text{Mn}(\text{CO})_4\text{Br}_2$  (d)  $\text{Mn}_2(\text{CO})_9\text{Br}$
31. The correct molecular representation of  $\text{W}(\text{Cp})_2(\text{CO})_2$  is, (Cp = cyclopentadienyl)  
(a)  $[\text{W}(\eta^1\text{-Cp})(\eta^3\text{-Cp})(\text{CO})_2]$  (b)  $[\text{W}(\eta^1\text{-Cp})(\eta^5\text{-Cp})(\text{CO})_2]$   
(c)  $[\text{W}(\eta^3\text{-Cp})(\eta^5\text{-Cp})(\text{CO})_2]$  (d)  $[\text{W}(\eta^5\text{-Cp})_2(\text{CO})_2]$
32. Match the metalloproteins with their respective functions.
- |                       |                         |
|-----------------------|-------------------------|
| P. Ferritin           | I. Electron transfer    |
| Q. Rubredoxin         | II. Acid-base catalysis |
| R. Cobalamin          | III. Metal storage      |
| S. Carbonic anhydrase | IV. Methyl transfer     |





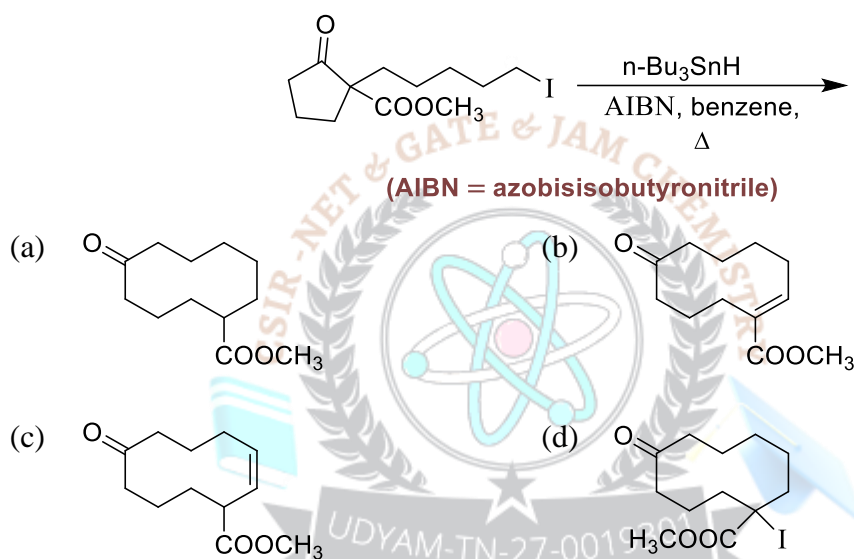
	P	Q	R	S
(a)	III ; II ; I ; IV			
(c)	IV ; I ; III ; II			

	P	Q	R	S
(b)	III ; I ; IV ; II			
(d)	IV ; II ; I ; III			

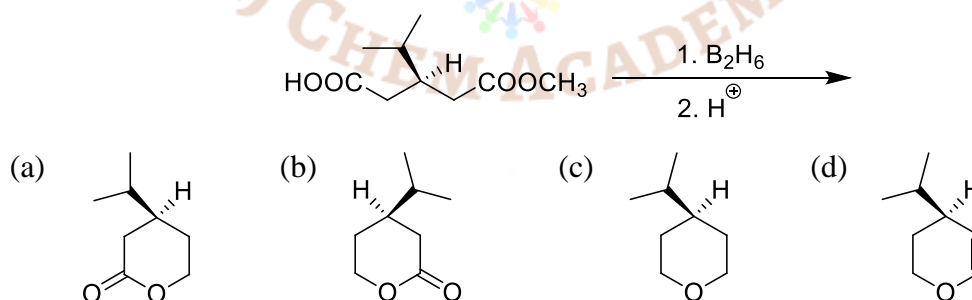
33. Suppose the wave function of a **one-dimensional** system is  $\varphi = \sin(kx) \exp(3ikx)$ . In an experiment measuring the momentum of the system, one of the **expected outcomes** is

- (a) 0 (b)  $\hbar k$  (c)  $2 \hbar k$  (d)  $3 \hbar k$

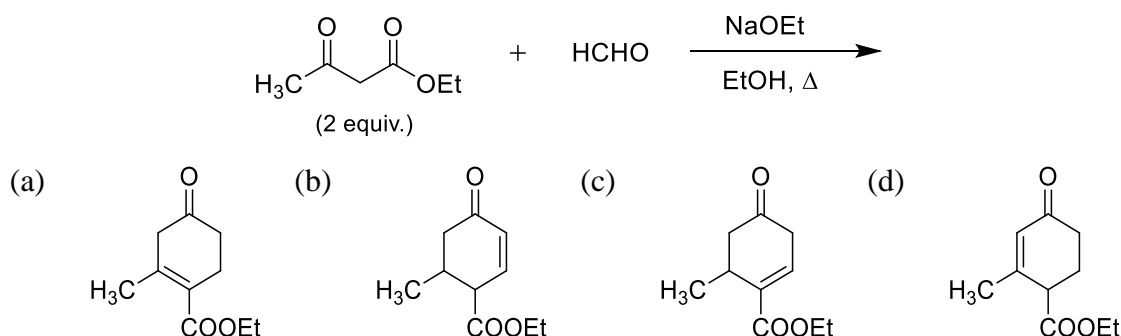
34. The **major product** formed in the following reaction is



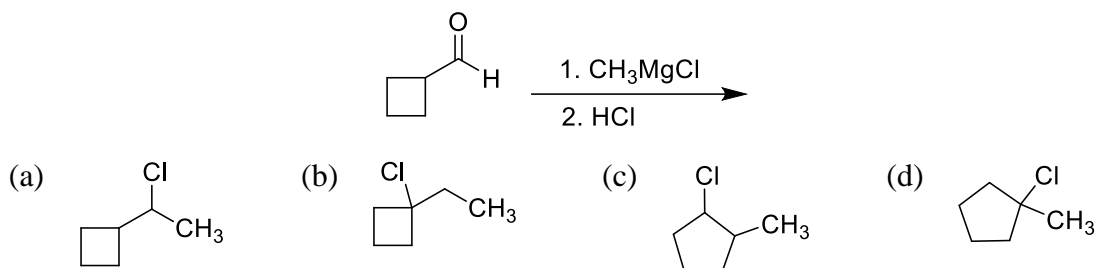
35. The **major product** formed in the following reaction is,



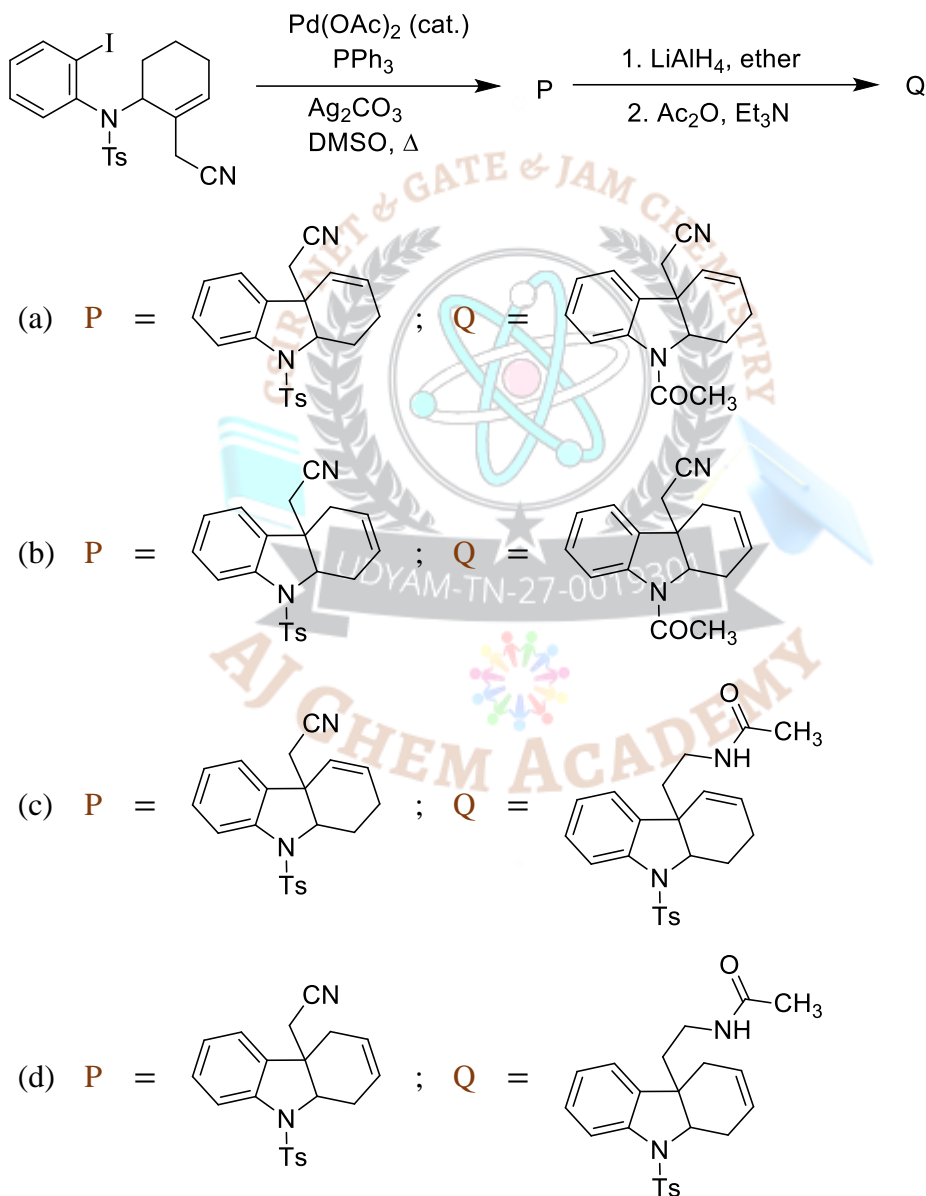
36. The **major product** formed in the following reaction is



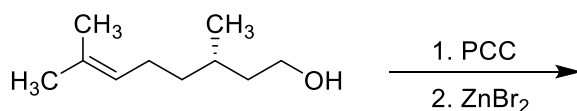
37. The **major product** formed in the following reaction is



38. In the following reaction sequence, the **products P and Q** are

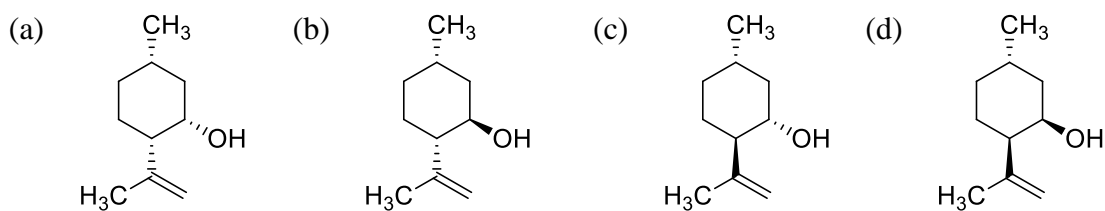


39. The **major product** formed in the following reaction is

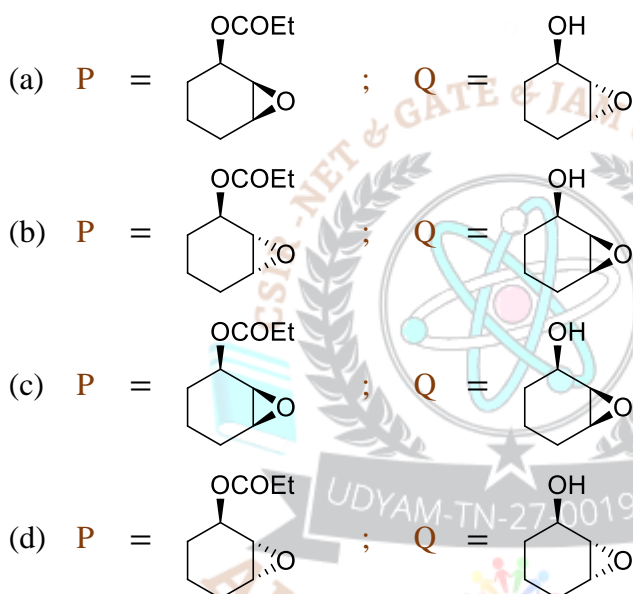
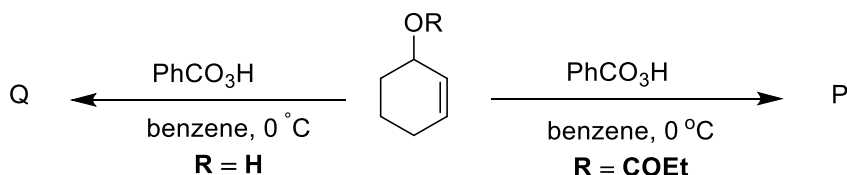


(PCC = Pyridinium chlorochromate)

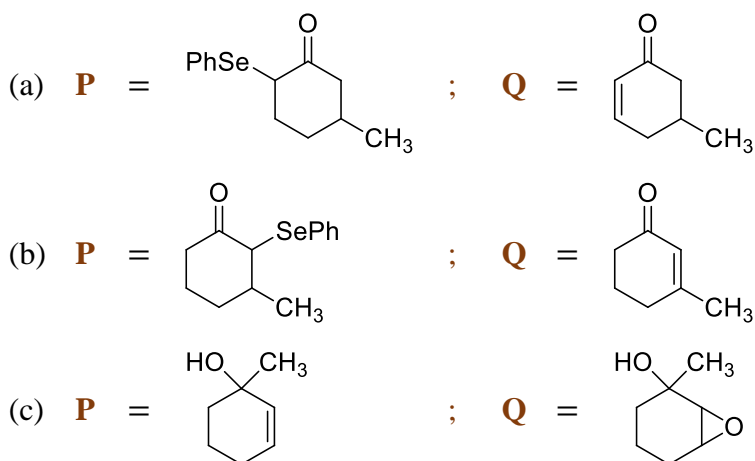
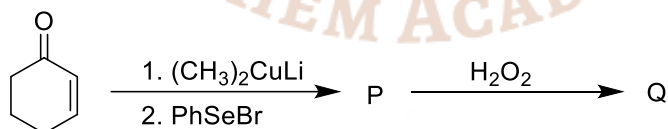


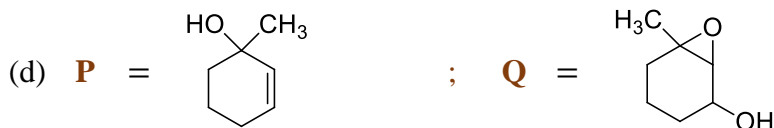


40. In the following reactions, the **major products P and Q** are

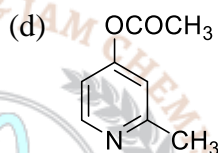
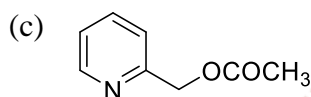
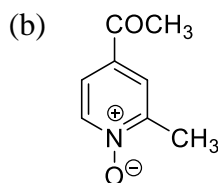
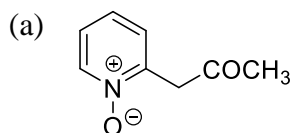
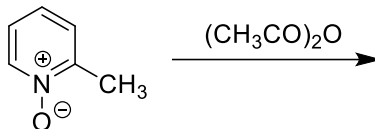


41. In the following reaction sequence, the **products P and Q** are





42. The **major product** formed in the following reaction is



43. The **rate** of the following redox reaction is **slowest** when **X** is,



(a)  $\text{H}_2\text{O}$

(b)  $\text{NH}_3$

(c)  $\text{Cl}^-$

(d)  $\text{N}_3^-$

44. A complex is composed of **one chromium ion, three bromides and six water molecules**. Upon addition of excess  $\text{AgNO}_3$ , 1.0 g aqueous solution of the complex gave 0.94 g of  $\text{AgBr}$ . The **molecular formula of the complex** is,

(Atomic weight:  $\text{Cr} = 52$ ,  $\text{Br} = 80$ ,  $\text{Ag} = 108$ ,  $\text{O} = 16$  and  $\text{H} = 1$ )

(a)  $[\text{Cr}(\text{H}_2\text{O})_6]\text{Br}_3$

(b)  $[\text{Cr}(\text{H}_2\text{O})_5\text{Br}]\text{Br}_2 \cdot \text{H}_2\text{O}$

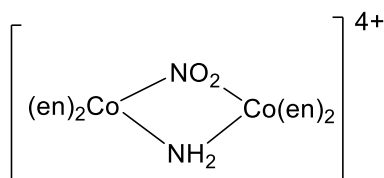
(c)  $[\text{Cr}(\text{H}_2\text{O})_4\text{Br}_2]\text{Br} \cdot 2\text{H}_2\text{O}$

(d)  $[\text{Cr}(\text{H}_2\text{O})_3\text{Br}_3] \cdot 3\text{H}_2\text{O}$

**Q.45 – Q.55 Numerical Answer Type (NAT), carry TWO marks each (no negative marks).**

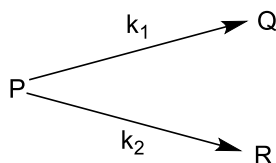
45. The **number of possible optically active isomer(s)** for the following complex is \_\_\_\_.

(en = ethylenediamine)



46. The **specific rotation** of optically pure (R)-2-bromobutane is  $-112.00$ . A given sample of 2-bromobutane exhibited a **specific rotation** of  $-82.88$ . The **percentage of (S)-(+)-enantiomer** present in this sample is \_\_\_\_\_.

47. Consider the following two parallel irreversible first order reactions at temperature T,



where  $k_1$  and  $k_2$  are the rate constants and their values are  $5 \times 10^{-2}$  and  $15 \times 10^{-2} \text{ min}^{-1}$ , respectively, at temperature T. If the initial concentration of the reactant 'P' is  $4 \text{ mol L}^{-1}$ , then the concentration of product 'R' after 10 min of reaction is \_\_\_\_\_  $\text{mol L}^{-1}$ . (Round off to two decimal places)

(Assume only P is present at the beginning of the reaction)

48. Consider the following equilibrium



At 298 K, the standard molar Gibbs energies of formation,  $\Delta_f G^0$ , of  $\text{SO}_{2(g)}$  and  $\text{SO}_{3(g)}$  are  $-300$  and  $-371 \text{ kJ mol}^{-1}$ , respectively. The value of the equilibrium constant,  $K_p$ , at this temperature is \_\_\_\_\_  $\times 10^{10}$ .

(Round off to the nearest integer & Gas constant  $R = 8.31 \text{ J mol}^{-1} \text{ K}^{-1}$ )

49. Consider the electrochemical cell:  $\text{M}_{(s)} | \text{MI}_{2(s)} | \text{MI}_{2(aq)} | \text{M}_{(s)}$ .

where 'M' is a metal. At 298 K, the standard reduction potentials are  $E_{\text{M}^{2+}(\text{aq})/\text{M}_{(s)}}^0 = -0.12 \text{ V}$ ,  $E_{\text{MI}_{2(s)}/\text{M}_{(s)}}^0 = -0.36 \text{ V}$  and the temperature coefficient is

$\left( \frac{\partial E_{\text{cell}}^0}{\partial T} \right)_P = 1.5 \times 10^{-4} \text{ VK}^{-1}$ . At this temperature the standard enthalpy change for

the overall cell reaction,  $\Delta_r H^0$ , is \_\_\_\_\_  $\text{kJ mol}^{-1}$ .

(Round off to two decimal places & Faraday constant  $F = 96500 \text{ C mol}^{-1}$ )

50. The normal boiling point of a compound (X) is 350 K (heat of vaporization,  $\Delta_{\text{vap}} H = 30 \text{ kJ mol}^{-1}$ ). The pressure required to boil 'X' at 300 K is \_\_\_\_\_ Torr.

(Round off to two decimal places & Ignore the temperature variation of  $\Delta_{\text{vap}} H$ )

(Gas constant  $R = 8.31 \text{ J mol}^{-1} \text{ K}^{-1}$  and  $1 \text{ atm} = 760 \text{ Torr}$ )

51. For a bimolecular gas phase reaction  $\text{P} + \text{Q} \rightarrow \text{R}$ , the pre-exponential factor is  $1 \times 10^{13} \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$ . The standard entropy of activation at  $25^\circ \text{C}$  is \_\_\_\_\_  $\text{JK}^{-1} \text{ mol}^{-1}$ . (Round off to two decimal points)

(The standard concentration  $c^0 = 1 \text{ mol dm}^{-3}$ ; Planck constant  $h = 6.62 \times$



$10^{-34}$  Js; Boltzmann constant  $k_B = 1.38 \times 10^{-23} \text{ J K}^{-1}$ ; Gas constant  $R = 8.31 \text{ J mol}^{-1} \text{ K}^{-1}$ )

52. Character table of point group  $D_8$  is given below.

$D_8$	E	$2C_8$	$2C_4$	$2C_8^3$	$C_2$	$4C_2'$	$4C_2''$
$A_1$	a	1	1	1	1	1	1
$A_2$	b	1	1	1	1	h	i
$B_1$	c	-1	1	-1	1	1	j
$B_2$	d	-1	1	-1	1	-1	1
$E_1$	e	$\sqrt{2}$	0	$-\sqrt{2}$	-2	0	0
$E_2$	f	0	-2	0	k	0	0
$E_3$	g	$-\sqrt{2}$	0	$\sqrt{2}$	-2	0	0

Value of  $(a + b + c + d + e + f + g + h + i + j + k)$  is equal to \_\_\_\_\_.

53. If  $\langle \alpha | \widehat{S}_x \widehat{S}_y - \widehat{S}_y \widehat{S}_x | \alpha \rangle = -i\hbar^2 a$  where  $\widehat{S}_x$  and  $\widehat{S}_y$  are spin angular momentum operators and  $|\alpha\rangle$  is spin up eigen function, then the value of 'a' is \_\_\_\_\_.

(Round off to one decimal place)

54. A particle in one dimensional box of length  $2a$  with potential energy

$$V = \begin{cases} 0 & |x| < a \\ \infty & |x| > a \end{cases}$$

is perturbed by the potential  $V' = cx \text{ eV}$ , where  $c$  is a constant. The 1<sup>st</sup> order correction to the 1<sup>st</sup> excited state of the system is \_\_\_\_\_  $\times c \text{ eV}$ .

55. Consider a two-dimensional harmonic oscillator with angular frequency  $\omega_x = 2\omega_y = 6.5 \times 10^{14} \text{ rad s}^{-1}$ . The wavelength of x polarized light required for the excitation of a particle from its ground state to the next allowed excited state is \_\_\_\_\_  $\times 10^{-6} \text{ m}$ . (Round off to one decimal place)

(Speed of light  $c = 3.0 \times 10^8 \text{ m s}^{-1}$ )

### Answer Key

Q.No	Ans		Q.No	Ans		Q.No	Ans
1.	d		21.	5.40 to 5.55		41.	b
2.	b		22.	1.25 to 1.30		42.	c
3.	d		23.	6		43.	b
4.	a		24.	3.9		44.	b
5.	b		25.	74 to 76		45.	2

## GATE – 2019 – CY

6.	b		26.	a		46.	13
7.	b		27.	b		47.	2.50 to 2.65
8.	a		28.	a		48.	265 to 295
9.	b		29.	d		49.	-38 to -37
10.	d		30.	a		50.	135 to 137
11.	b		31.	c		51.	-12.9 to -12.4
12.	b		32.	b		52.	9
13.	c		33.	c		53.	0.5
14.	4		34.	a		54.	0
15.	5		35.	b		55.	2.8 to 3.0
16.	5		36.	d			
17.	3		37.	d			
18.	5		38.	c			
19.	11		39.	c			
20.	-126 to -120		40.	b			

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