Test Booklet Code & No. प्रश्नपत्रिका कोड व क्र.

В

Paper-III

AL SCIENCE
Seat No.
(In figures as in Admit Card)
Seat No.
(In words)
OMR Sheet No.
(To be filled by the Candidate)
[Maximum Marks : 150
Number of Questions in this Booklet: 75
विद्यार्थ्यांसाठी महत्त्वाच्या सूचना 1. परिक्षार्थींनी आपला आसन क्रमांक या पृष्ठावरील वरच्या कोप-यात लिहावा. तसेच आपणांस दिलेल्या उत्तरपत्रिकंचा क्रमांक त्याखाली लिहावा. 2. सदर प्रश्नपत्रिकंत 75 बहुपर्यायी प्रश्न आहेत. प्रत्येक प्रश्नास दोन गुण आहेत. या प्रश्नपत्रिकंतील सर्व प्रश्न सोडिवणे अनिवार्य आहे. सदरचे प्रश्न हे या विषयाच्या संपूर्ण अभ्यासक्रमावर आधारित आहेत. 3. परीक्षा सुरू झाल्यावर विद्यार्थ्याला प्रश्नपत्रिका दिली जाईल. सुरुवातीच्या 5 मिनीटांमध्ये आपण सदर प्रश्नपत्रिका उघडून खालील बाबी अवश्य तपासून पहाव्यात. (i) प्रश्नपत्रिका उघडण्यासाठी प्रश्नपत्रिकंवर लावलेले सील उघडावे. सील नसलेली किंवा सील उघडलेली प्रश्नपत्रिकं एकूण पृष्ठे तसेच प्रश्नपत्रिकंतील एकूण प्रश्नांची संख्या पडताळून पहावी. पृष्ठे कमी असलेली/कमी प्रश्न असलेली/प्रश्नांचा चूकीचा क्रम असलेली किंवा इतर त्रुटी असलेली सदोष प्रश्नपत्रिका सुश्वातीच्या 5 मिनिटातच पर्यवेक्षकाला परत देऊन दुसरी प्रश्नपत्रिका मागवून घ्यावी. त्यानंतर प्रश्नपत्रिका बदलून मिळणार नाही तसेच वेळही वाढवून मिळणार नाही याची कृपया विद्यार्थांनी नोंद घ्यावी. (iii) वरीलप्रमाणे सर्व पडताळून पहिल्यानंतरच प्रश्नपत्रिकंवर ओ.एम.आर. उत्तरपत्रिकंचा नंबर लिहावा. 4. प्रत्येक प्रश्नासाठी (A), (B), (C) आणि (D) अशी चार विकल्प उत्तरे दिली आहेत. त्यातील योग्य उत्तराचा रकाना खाली दर्शविल्याप्रमाणे ठळकपणे काळा/निळ करावा.
उदा. : जर (C) हे योग्य उत्तर असेल तर. A B D
 ग्रा प्रश्नपत्रिकेतील प्रश्नांची उत्तरे ओ.एम.आर. उत्तरपत्रिकेतच दर्शवावीत. इतर ठिकाणी लिहीलेली उत्तरे तपासली जाणार नाहीत. आत दिलेल्या सूचना काळजीपूर्वक वाचाव्यात. प्रश्नपत्रिकेच्या शेवटी जोडलेल्या को-या पानावरच कच्चे काम करावे. जर आपण ओ.एम.आर. वर नमूद केलेल्या ठिकाणा व्यतिरीक्त इतर कोठेही नाल, आसन क्रमांक, फोन नंबर किंवा ओळख पटेल अशी कोणतीही खूण केलेली आढळून आल्यास अथवा असभ्य भाषेचा वापर किंवा इतर गैरमागांचा अवलंब केल्यास विद्यार्थ्याला परीक्षेस अपात्र ठरविण्यात येईल. परत करणे आवश्यक आहे. तथापी, प्रश्नपत्रिका व ओ.एम.आर. उत्तरपत्रिकेची द्वितीय प्रत आपल्याबरोबर नेण्यास विद्यार्थ्यांना परवानगी आहे. फक्त निळ्या किंवा काळ्या बॉल पेनचाच वापर करावा. कलक्युलेटर किंवा लॉग टेबल वापरण्यास परवानगी नाही. चुकीच्या उत्तरसाठी गुण कपात केली जाणार नाही.

11. 12.

MAY - 33316/III—B

Chemical Science Paper III

Time Allowed: 2½ Hours] [Maximum Marks: 150

Note: This paper contains **Seventy Five (75)** multiple choice questions, each question carrying **Two (2)** marks. Attempt *All* questions.

- 1. Which of the following statements is *not correct* about the quantum mechanical harmonic oscillator ?
 - (A) The amplitude of a quantum mechanical harmonic oscillator in its ground state will not exceed its classical value in lower energy states.
 - (B) The average values of $\langle x \rangle$ and $\langle p \rangle$ are zero.
 - (C) In highly excited states the quantum mechanical oscillator behaves as a classical oscillator
 - (D) The wave function for quantum mechanical harmonic oscillator is always either odd or even.
- 2. The free energy of mixing for the ideal solution is given by :

$$\Delta G_{mix} = RT(x_A \ln x_A + x_B \ln x_B)$$

 $\because x_{A}$ and x_{B} are respective mole fractions from the expression one can deduce that :

- (A) The mixing is not spontaneous and $\Delta G = +ve$
- (B) The mixing is spontaneous and ΔG is negative and maximum at X=1/2
- (C) The mixing is exothermic
- (D) The mixing is endothermic

- 3. The term symbol not possible for the nd^2 electronic configuration is :
 - (A) ^{3}F

(B) ^{1}D

(C) ${}^{1}G$

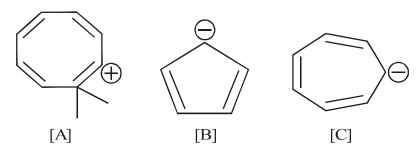
- (D) ^{1}F
- 4. Log of mean activity coefficient (γ_{\pm}) for 0.005 mol kg^{-1} KCl solution at $25^{\circ}C$ is :
 - (A) -0.920

(B) 0.0360

(C) 0.920

(D) -0.0360

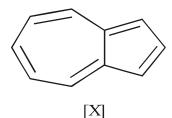
5.



Among the ions [A], [B] and [C], given:

- (A) A is aromatic, B is homoaromatic and C is antiaromatic
- (B) A is homoaromatic, B is aromatic and C is antiaromatic
- (C) A is aromatic, B is antiaromatic and C is homoaromatic
- (D) A is homoaromatic, B is antiaromatic and C is aromatic

6.



The most correct statement about compound [X] is :

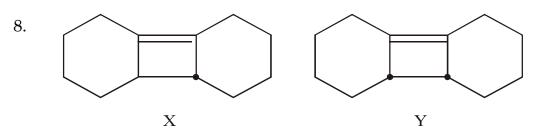
- (A) It is aromatic
- (B) It is aromatic and has high dipole moment than expected
- (C) It is aromatic but has no dipole moment
- (D) It is antiaromatic

7. Justify the answer given in the statement below for reaction (X) and reaction (Y) pathways:

$$Br_2 + \ddot{S} \underbrace{\stackrel{CH_3}{\leftarrow}}_{CH_3} \underbrace{\stackrel{Solvent}{\rightarrow}}_{Catalyst} Br + Br \underbrace{\stackrel{\bigoplus}{\leftarrow}}_{S} \underbrace{\stackrel{CH_3}{\leftarrow}}_{CH_3} (X) Reaction Path$$

$$CH_3$$
— CH_3 + \ddot{S}
 CH_3
 CH_3
 CH_3
 CH_3
 CH_3
 CH_3
 $Colvent$
 $Colven$
 $Colvent$
 $Colven$
 $Colvent$
 $Colvent$
 $Colvent$
 $Colven$
 Co

- (A) Lone pair of electrons are donated from S-atom to Br—Br to form new S-Br band, in (X) path but not possible to break C—C bond in (Y) path
- (B) Lone pair of electrons are donated from Br—Br to S-atom in path (X), but not possible to donate CH_3 — CH_3 σ -bond to S-atom in path (Y)
- (C) Lone pair of electrons are restricted by S-atom to donate in both (X) path and (Y) path, but catalytic reagent may do this in path (X) only
- (D) Lone pair of electrons are blocked by S-atom in path (Y) but bromine procure electron pair from solvent in path (X)



Both X and Y are subjected to thermal ring opening. It will be observed that:

- (A) X undergoes ring opening, more readily compared to Y
- (B) Y undergoes ring opening, more readily compared to X
- (D) Y does not undergo ring opening even at high temperature

9. Methoxymethyl ether of *m*-cresol is reacted with *t*-BuLi in dry ether and the product is reacted with dry ice followed by acidic work-up. The major product (X) formed has structure:

The above reaction involves:

- (A) [1, 3] sigmatropic carbon migration with inversion at migrating center
- (B) [1, 3] sigmatropic carbon migration with retention at migrating center
- (C) [1, 3] sigmatropic hydrogen migration
- (D) [1, 5] sigmatropic carbon migration with retention at migrating center

- 11. Pericyclic disrotatory ring opening of 1, 3-cyclo hexadiene to 1, 3, 5- hexatriene could be considered as:
 - (A)
- $[\pi^2 s + \sigma^2 a]$ cycloaddition (B) $[\pi^2 a + \sigma^2 a]$ cycloaddition
 - (C)
- $[\pi^4 s + \sigma^2 s]$ cycloaddition (D) $[\pi^4 a + \sigma^2 a]$ cycloaddition
- An aromatic ether (C_7H_8O) is reacted with n-butyl lithium and the ether 12. derivative obtained is reacted with an aliphatic aldehyde to form $Y(C_9H_{12}O_2)$. Y on oxidation forms compound $Z(C_9H_{10}O_2)$. Compound Z has the following PMR data:

The structure of compound [Y] is:

MAY - 33316/III—B

Compounds (X) and (Y) exhibit two singlets each in their ¹H NMR spectrum.

The expected chemical shifts in δ units are :

- (A) 2.3 and 7.2 for X; 0.95 and 4.8 for Y
- (B) 0.95 and 4.8 for X; 2.3 and 7.2 for Y
- (C) 0.95 and 7.2 for X; 2.3 and 4.8 for Y
- (D) 2.3 and 4.8 for X; 0.95 and 7.2 for Y
- 14. Identify the name of product (X) in the following transformation :

$$HO \longrightarrow OH$$

$$(X)$$

(A) X = Ketal

(B) X = Lactal

(C) X = Acetal

(D) X = Hemiketal

15. The major product (X) in the following reaction is :

$$\begin{array}{c|c}
\hline
N & DMF \\
\hline
POCl_3
\end{array}$$

$$(A) \qquad \overbrace{\hspace{1cm} N \hspace{1cm}} CHO$$

(C)
$$V$$

$$(D) \qquad \begin{array}{c} \text{CHO} \\ \text{N} \\ \text{H} \end{array}$$

16. In the following reaction, first step involved deprotonation of chloroform by strong base (usually KOH) to form the reactive intermediate (X) as:

OH
$$+ CHCl_{3} + KOH_{(aq)} \xrightarrow{Shaking} [X]$$
(Reactive Intermediate)

$$\xrightarrow{\text{Step (II)}} \text{CHO} \\ + \text{H}_2\text{O Main product}$$

- (A) X = Dichlorocarbanion ion
- (B) X = Trichlorocarbanion ion
- (C) X = Dichlorocarbene species
- (D) X = Chlorohydrin species

The above reaction is an example of reaction and structure of $\left[X\right]$ is :

- (A) Stevens rearrangement, X =
- (B) Shapiro reaction, X =
- (C) Shapiro reaction, X =
- (D) Stevens rearrangement, X =

18. Identify the following conversion:

- (A) Ene Reaction
- (B) Hofmann Loffler Reaction
- (C) Barton Reaction
- (D) Shapiro Reaction

19.	Thyn	nine is a pyrimidine base present in DNA, its carbohydrate substitution			
	in 2-	2-deoxyribofuranose is attached to :			
	(A)	N-3 of Thymine			
	(B)	N-1 of Thymine			
	(C)	Both N-1 and N-3 of Thymine			
	(D) Neither N-1 nor N-3 of Thymine				
20.	Mate	atch the following combinations and choose suitable pairs :			pose suitable pairs :
	(L)	β-Carotene		(1)	Protein
	(M)	Morphine		(2)	Alkaloid
	(N)	Alanin		(3)	Steroid
	(O)	Estrogene		(4)	Terpene
	(A)	(L)—(3)	(M)—(1)	(N)—(4)	(O)—(2)
	(B)	(L)—(1)	(M)—(3)	(N)—(2)	(O)—(4)
	(C)	(L)—(4)	(M)—(2)	(N)—(1)	(O)—(3)
	(D)	(L)—(2)	(M)—(4)	(N)—(3)	(O)—(1)

- 21. One of the statement is true in case of Nucleotides:
 - (A) They are phosphoric acid esters of Pyrimidines
 - (B) They are phosphoric acid esters of Sugars
 - (C) They are phosphoric acid esters of Purines
 - (D) They are phosphoric acid esters of Nucleosides
- 22. The oil of winter green (X) is formed in one of the following reactions:

(A)
$$COOH$$
 + CH_3COOH $Refluxed$ (X)

(B)
$$+ \text{HCOOH} \xrightarrow{\text{Refluxed}} \text{(X)}$$

(C) OH +
$$CH_3COOH \xrightarrow{H_2SO_4} (X)$$

(D)
$$COOH + CH_3OH \xrightarrow{H_2SO_4} (X)$$

- 23. The carbonyl compounds exhibits electrophilic functionality because they have :
 - (A) A Low Energy LUMO σ* orbital
 - (B) A Low Energy HOMO π^* orbital
 - (C) A Low Energy LUMO π^* orbital
 - (D) A Low Energy HOMO σ^* orbital
- 24. In emulsion polymerization method, the initiator used is:
 - (A) Soluble in monomer
 - (B) Soluble in water
 - (C) Soluble in both water and monomer
 - (D) Insoluble in both water and monomer
- 25. Simple following alkene is a weak nucleophile and reacts with X = electrophile given below, identify the best suitable electrophile and Y = product :

Alkene

Product

- (A) $X = Br_2$ and Y = Dibromoalkane
- (B) X = NO and Y = Nitrosoalkane
- (C) $X = NO_2$ and Y = Nitroalkane
- (D) X = CO and Y = Carboxylalkane
- 26. ${}^{\Theta}BH_4$ can have electrons from B-H bond available for donation, although ${}^{\Theta}BH_4$ devoid lone pair of electrons of π orbital electron it act as:
 - (A) poor nucleophile

(B) good nucleophile

(C) good electrophile

(D) neutral species

Linked problems, Q. No. 27 and Q. No. 28:

27. The structure of [X] is:

28.	The name of the above reaction is:				
	(A)	Paterno-Buchi reaction	(B)	Barton reaction	
	(C)	Michael addition	(D)	Ene reaction	
29.	9. When gamma-ray photon with sufficiently high energy interacts with ma			igh energy interacts with matter,	
	pair production can occur. In this process, the photon is totally absorbed				
	creat	ing:			
	(A)	a positron	(B)	an electron	
	(C)	a positron and an electron	(D)	a neutron	
30.	Two 10.0 ml aliquots of sea water were taken for neutron activation analys			en for neutron activation analysis.	
	Exactly 1.0 ml of a standard solution containing 1.0 μg Al^{3+} was added to				
	one aliquot and 1.0 ml of distilled water was added to other aliquot. The two				
	samples were then irradiated simultaneously in a homogeneous neutron flux.				
	After a brief cooling period, the gamma radiation from the decay of 28 Al was				
	counted for each sample. The solution diluted with distilled water gave a				
	counting rate of 2200 cpm whereas the solution containing the added Al ³⁺				
	gave 4200 cpm. What is the mass of Al in the 10 ml aliquot?				
	(A)	2.2 μg	(B)	1.1 μg	
	(C)	2.4 μg	(D)	1.2 μg	

- 31. Which one of the following is an example of β -decay?
 - (A) ${}^{48}_{24}\text{Cr} + {}^{0}_{1}\text{e} \rightarrow {}^{48}_{23}\text{V} + \text{X-rays}$
 - (B) $^{238}_{92}\text{U} \rightarrow ^{234}_{90}\text{Th} + ^{4}_{2}\text{He}$
 - (C) ${}^{9}_{4}\text{Be} + {}^{4}_{2}\text{He} \rightarrow {}^{12}_{6}\text{C} + {}^{1}_{0}\text{n} + 5.7\text{MeV}$
 - (D) ${}^{23}_{11}Na + {}^{1}_{0}n \rightarrow {}^{24}_{11}Na + gamma rays$
- 32. Edge dislocations result when:
 - (A) atoms in one layer do not match up precisely with those of the next
 - (B) atoms in one layer do match up with those of the next
 - (C) cations migrate to the surface
 - (D) anions migrate to the surface
- 33. Frenkel defect is caused when:
 - (A) Anions are missing from its lattice sites
 - (B) Cations are migrated to nearby interstitial sites
 - (C) Cations and equal number of anions migrate from its lattice sites to the surface of crystal
 - (D) Cations are migrating from its lattice sites to the surface of crystal
- 34. For transition metal (M) spinels, $M(II) [M'III]_2O_4$ is considered a normal spinel, while $M'(III) [M(II) M'(III)]O_4$ is considered an inverse spinel. Based on these consideration which one of the following is true?
 - (A) Fe₃O₄ is a normal spinel
 - $(B) \hspace{0.5cm} Ni(II)Fe_2O_4 \hspace{0.1cm} is \hspace{0.1cm} a \hspace{0.1cm} normal \hspace{0.1cm} spinel \\$
 - (C) Co₃O₄ is an inverse spinel
 - $(D) \hspace{0.5cm} Mn_3O_4 \hspace{0.1cm} is \hspace{0.1cm} a \hspace{0.1cm} normal \hspace{0.1cm} spinel \\$

		19	9	[P.T.O			
	(C)	Ru ³⁺	(D)	V^{3+}			
	(A)	Fe^{2+}	(B)	$\mathrm{Fe^{3+}}$			
	comp	plex with pyridine?					
38.	Whic	Which of the following metal ions should form a more stable octahedra					
	(D)	D) Broad and appear at lower wavelength					
	(C)	sharp and line-like					
	(B)	broad and high intensity					
	(A)	broad and low intensity					
37.	The	absorption bands in electronic	spectra	a of lanthanide cations are :			
		magnitude					
	(D)	the spin-orbital coupling and	d ligano	d field effects are of comparable			
	(C)	the 5f electrons interact much	ch lesse	er with ligands			
	(B)	the 5f electrons do not inter	act wit	h ligands			
	(A)	spin-orbital coupling of 5f el	ectrons	is negligible			
36.	The	The magnetic properties of the actinides are quite complex because :					
	(C)	3003	(D)	4012			
	(A)	2002	(B)	4120			
35.	The	correct STYX number for ${\rm B_5H_9}$ (nido-pentaborane) is :					

	(A)	$[\mathrm{Mn(CN)}_6]^{4-}$	(B)	$[Mn(CN)_6]^{3-}$
	(C)	$[{\rm Mn(H_2O)_6}]^{5+}$	(D)	$[\mathrm{FeF}_{6}]^{3-}$
40.	The	number of valence electrons in	Ni(η ^ξ	3 -allyl) $_{2}$ and $\mathrm{Mn}(\eta^{5}\text{-}\mathrm{C}_{5}\mathrm{H}_{5})$ (CO) $_{3}$
	is, re	espectively:		
	(A)	18 and 17	(B)	16 and 18
	(C)	14 and 19	(D)	16 and 17
41.	1. How many M-M bonds should form in complex $[Mo(\eta^5-C_5H_5) (CO)_2]$ order to obey 18 electron rule ? [Given : At. no. of Mo = 42]			
	(A)	1	(B)	2
	(C)	3	(D)	4
42.	Whic	h of the following is a stable 1	.8 elec	tron species ?
	(A)	$\mathrm{Ni}(\eta^3\text{-allyl})_2$		
	(B)	$Ru(\eta^{5}\text{-}{\rm C}_{7}{\rm H}_{7})\ (\eta^{1}\text{-}{\rm C}_{6}{\rm H}_{5}){\rm CO}$		
	(C)	${\rm Mo}(\eta^5\text{-}{\rm C}_5{\rm H}_5)\ (\eta^1\text{-}{\rm C}_5{\rm H}_5)\ ({\rm CO})_2$		
	(D)	$\mathrm{Rh}(\mathrm{PPh}_3)_2\mathrm{COCl}$		
43.	The π -acidic ligands stabilize the metal ions in lower oxidation states because of :			
	(A)	$L \rightarrow M$ π -interaction		
	(B)	Stronger L \rightarrow M σ -interaction	1	
	(C)	Synergic effect		
	(D)	Neutralization of the positive	charge	e over the metal ion.
		20		

Which of the following complexes has zero orbital moment?

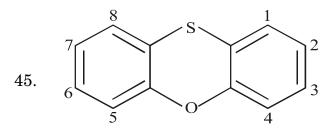
39.

- 44. Which of the following will undergo the easiest hydrogenation?
 - $(A) \qquad Cr(\eta^6\text{-}C_6H_6)_2$

 $(B) \qquad Fe(\eta^5\text{-}C_5H_5)_2$

 $\mathrm{(C)} \quad \ Mo(\eta^5\text{-}\mathrm{C_5H_5})_2$

(D) $Co(\eta^5 - C_5H_5)_2$



Yields a monolithiated product on lithiation with Li(s) in an inert solvent. Which C atom is expected to exchange H with Li?

(A) 1

(B) 2

(C) 3

- (D) 4
- 46. During a ligand substitution reaction, if an intermediate with higher coordination is formed, the possible mechanism of the reaction is:
 - (A) A

(B) I_a

(C) D

- (D) I_d
- 47. Which of the following is expected to undergo the fastest inner sphere reduction by $[Cr(H_2O)_6]^{2+}$?
 - $(A) \qquad [\mathrm{Co(PY)}_{6}]^{3+}$

- $(B) \quad [\text{Co(PY)}_5 \ (\text{ampy})]^{3+}$
- $(C) \quad [Fe(PY)_5 \ (dampy)]^{3+}$
- (D) $[Co(PY)_5 (dampy)]^{3+}$

[PY = pyridine

ampy = 4-aminopyridine

dampy = 3, 5-diaminopyridine]

48. The reaction:

$$\left[Pt(en)_{2}Cl_{2} \right]^{2+}_{(aq)} + \left[Pt(NH_{3})_{4} \right]^{2+}_{(aq)}$$

$$\xrightarrow{Cl^{-}} \left[Pt(en)_{2} \right]^{2+}_{(aq)} + \left[Pt(NH_{3})_{4}Cl_{2} \right]^{2+}_{(aq)}$$

is an example of:

- (A) Isomerization reaction
- (B) Ligand substitution reaction
- (C) Electron transfer process
- (D) Reproportionation reaction

49. A 4Fe-3S protein has:

- (A) 4 sulphides and 3Fe bound to cysteine residues
- (B) 3 sulphides and 4Fe bound to cysteine residues
- (C) 4 Fe and 3 cysteine residues
- (D) 4 sulphides out of which 3 are bridging between the Fe ions

50. The MET-APO form of the active site in hemocyanin has:

(A) One Cu⁺

- (B) One Cu^{2+}
- (C) One Cu^{1+} and one Cu^{2+}
- (D) Two Cu^{2+}

51. Which of the following has the lowest nephelauxetic ratio?

 $(A) [RuCl_6]^{3-}$

(B) $[Fe(PY)_4Cl_2]$

(C) $[Ru(NH_3)_6]^{2+}$

(D) $[Ru(bipy)_3]^{2+}$

[bipy = 2, 2'-bipyridine]

[PY = pyridine]

52. On the basis of cluster valence electron count, the structure of the metallic framework in ${\rm Co_4(CO)_{12}}$ and ${\rm Ru_4(CO)_{16}}$ can be predicted to be respectively :

- (A) Tetrahedral and square planar
- (B) Square planar and butterfly
- (C) Butterfly and tetrahedral
- (D) Square planar and tetrahedral

53. A solution containing 10.0 ppm $\rm KMnO_4$ had a transmittance of 0.1 in a 2.0 cm cell at 520 nm. The absorptivity of $\rm KMnO_4$ solution is :

- (A) $5.0 \times 10^2 \text{ Lg}^{-1} \text{ cm}^{-1}$
- (B) $5.0 \times 10^3 \text{ Lg}^{-1} \text{ cm}^{-1}$
- (C) $2.5 \times 10^2 \text{ Lg}^{-1} \text{ cm}^{-1}$
- (D) $2.5 \times 10^3 \text{ Lg}^{-1} \text{ cm}^{-1}$

54. An acid HB has a distribution coefficient of 8.8 between an organic solvent and water. At pH 6.0, exactly half of the material is extracted into the organic solvent from an equal volume of the aqueous phase. The dissociation constant of acid HB is:

(A) 8.8×10^{-6}

(B) 7.8×10^{-6}

(C) 9.8×10^{-6}

(D) 6.8×10^{-6}

55. Which of the following is used as sources of radiation in uv-visible spectrophotometer?

- (A) Tungsten lamp
- (B) Deuterium lamp
- (C) Tungsten and deuterium lamp
- (D) Xenon arc lamp

- 56. If the vivrational frequency of H_2 molecule is 4400 cm⁻¹, what will be the frequency of vibration of HD?
 - (A) 2200 cm^{-1}

(B) 4400 cm^{-1}

(C) 3300 cm^{-1}

- (D) 3833 cm^{-1}
- 57. The electrochemical cells are connected as shown below:

$$\begin{array}{c|c} Zn(s) & Zn^{2+}(1.0M) \, \| \, Cu^{2+}(0.1M) | \, \, Cu(s) \, \, cell \, \, I \\ & | & | \\ Zn(s) & | \, \, Zn^{2+}(0.1M) \, \| \, Cu^{2+}(1.0M) | \, \, Cu(s) \, \, cell \, \, II \end{array}$$

Which statement is *correct*?

- (A) Flow of electrons from zinc electrode of cell (II) to the zinc electrode of cell (I)
- (B) No current will flow
- (C) Flow of electron from zinc electrode of cell (I) to the zinc electrode of cell (II)
- (D) $E_{\text{cell}} = E_{\text{Cell}}^0$
- 58. The free energy change of 1 mole of an ideal gas compressed isothermally from 1 atm to 2 atm is:
 - (A) -2 RT

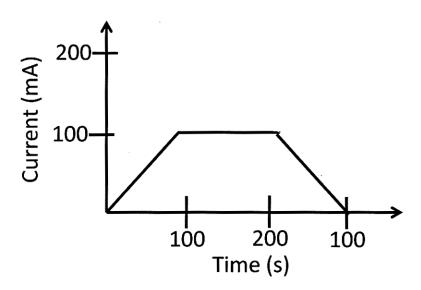
(B) RT ln 2

(C) -RT ln 2

(D) 2 RT

59. The following graph represents current-time curve for the deposition of Cu.

Based on this graph, the amount of Cu deposited would be (Given : Atomic weight of Cu = 63) :



(A) 6.3 g

(B) 63 g

(C) $6.3 \times 10^{-3} \text{ g}$

- (D) $6.3 \times 10^{-4} \text{ g}$
- 60. Which of the following shows maximum osmotic pressure in water?
 - (A) 1 M NaCl

- (B) 1 M MgCl₂
- (C) 1 M (NH₃)₃ PO₄
- (D) $1 \text{ M Na}_2\text{S}_2\text{O}_4$
- 61. Which of the following reactions is involved in the fuel cell?
 - ${\rm (A)} \hspace{0.5cm} {\rm Cd} \hspace{0.1cm} + \hspace{0.1cm} 2{\rm Ni(OH)}_{3} \hspace{0.1cm} \rightarrow \hspace{0.1cm} {\rm CdO} \hspace{0.1cm} + \hspace{0.1cm} 2{\rm Ni(OH)}_{2} \hspace{0.1cm} + \hspace{0.1cm} {\rm H}_{2}{\rm O}$
 - (B) Pb + PbO₂ + 2H₂SO₄ \rightarrow 2PbSO₄ + 2H₂O
 - ${\rm (C)} \hspace{0.5cm} 2{\rm H}_2 \hspace{0.1cm} + \hspace{0.1cm} {\rm O}_2 \hspace{0.1cm} \rightarrow \hspace{0.1cm} 2{\rm H}_2{\rm O}$
 - (D) 2Fe + O₂ + 4H⁺ \rightarrow 2Fe²⁺ + 2H₂O

62. The trial function for the particle in 1D box (with $0 \le x \le a$) leading to the best possible ground state energy employing the variational method should be:

(A) $x^{\alpha}(a-x)^{\beta}$

(B) $\sin\left(\frac{\pi x}{a}\right)$

(C) x(a-x)

(D) $x^2(a-x)^2$

63. 24 J is consumed to blow a soap bubble of the radius 1 cm. The surface tension at air-water interface would be around :

(Hint : W = $\gamma \times A$)

(A) 100 Jm^{-2}

(B) 10 Jm^{-2}

(C) 1 Jm⁻²

(D) 1000 Jm^{-2}

64. Out of the following which of the point group can a molecule have a dipole moment?

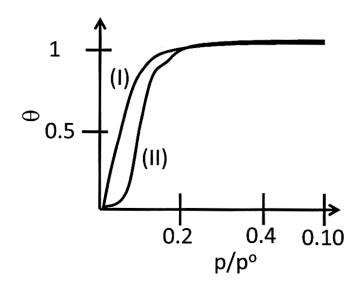
(A) C_{2y}

(B) D_{2d}

(C) D_{2h}

 $(D) \quad \ \, C_{\infty h}$

65. Which statement is appropriate for the following two adsorption isotherms?



- (A) Plot (I) is a classic case of lateral interaction i.e. Frumkin-Fowler-Guggenheim (FFG) adsorption isotherm
- (B) Plot (II) is classic case of Langmuir adsorption isotherm
- (C) Plot (II) is a case of FFG adsorption isotherm
- (D) Practically there is no difference between (I) and (II)
- 66. Which of the following relations is not correct for an ideal gas?

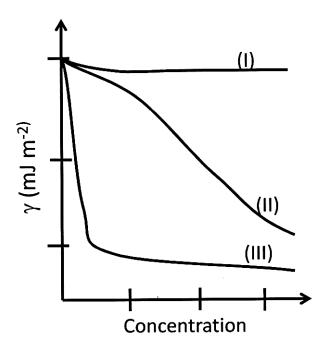
(A)
$$\left(\frac{\partial H}{\partial T}\right)_p - R = \left(\frac{\partial H}{\partial T}\right)_v$$

(B)
$$\left(\frac{\partial H}{\partial T}\right)_p > \left(\frac{\partial H}{\partial T}\right)_v$$

(C)
$$\left(\frac{\partial U}{\partial V}\right)_T = 0$$

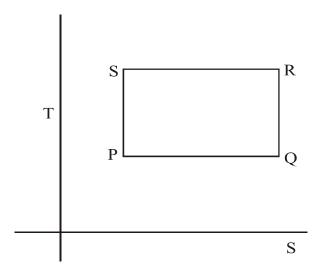
(D)
$$\left(\frac{\partial H}{\partial T}\right)_{p} + R = \left(\frac{\partial H}{\partial T}\right)_{v}$$

67. Consider the following three plots for the surface tensions noted for three different solutes. Based on this which statement is most appropriate?



- (A) All three plots are for surface active agents.
- (B) From these plots, the critical micelles concentrations can be estimated.
- (C) Among these, plot (III) is for a surfactant and plot (I) is possibly for an electrolyte.
- (D) Among (II) and (III), plot II is for more active surface active agent.

68. If the Carnot cycle in enthalpy-temperature diagram shows:



then the system rejects heat to the surroundings in going from:

$$(A)$$
 $R \rightarrow S$

$$(B) \quad P \rightarrow Q$$

$$(C)$$
 $Q \rightarrow P$

$$(D)$$
 $S \rightarrow R$

69. In the Michaelis-Menten mechanism for the enzymolysis:

$$\mathbf{E} + \mathbf{P} \xrightarrow{K_1} \mathbf{X} \xrightarrow{K_2} \mathbf{E} + \mathbf{P}$$

The steady-state approximation leads to a concentration of an intermediate:

$$[X] = \frac{K_1 E_0 S_0}{K_1 S_0 + K_{-1} + K_2}$$

29

Then the initial rate of reaction is given as:

(A)
$$v_0 = \frac{K_2 S_0}{1 + \left(\frac{K_{-1} + K_2}{K_1}\right) \frac{1}{E_0}}$$
 (B) $v_0 = \frac{V_m}{1 + \frac{K_m}{E_0 S_0}}$

(B)
$$v_0 = \frac{V_m}{1 + \frac{K_m}{E_0 S_0}}$$

(C)
$$v_0 = \frac{K_2 E_0}{1 + \left(\frac{K_{-1} + K_2}{K_1}\right) \frac{1}{S_0}}$$
 (D) $v_0 = \frac{V_m}{1 + \frac{K_m}{E_0}}$

$$(\mathbf{D}) \qquad \mathbf{v}_0 = \frac{\mathbf{V}_m}{1 + \frac{\mathbf{K}_m}{\mathbf{E}_0}}$$

70. What is the de Broglie wavelength associated with a particle in 1D box $(0 \le x \le a)$ with the width of 5 Å in its first excited state (the PE inside the box is zero and that outside being infinity):

(A) 10 Å

(B) 5 Å

(C) 2.5 Å

(D) 1 Å

71. Laughing gas N_2O , can be prepared from H_2 and NO:

$${\rm H_2(g)} \, + \, 2{\rm NO(g)} \, \to \, {\rm N_2O(g)} \, + \, {\rm H_2O(g)},$$

A study of initial concentration verses initial rates at certain tempeature yield the following data:

Set	[H ₂], M	[NO], M	$ m R_0~Ms^{-1}$
1	0.1	0.5	2.560×10^{-6}
2	0.2	0.3	1.843×10^{-6}
3	0.1	0.3	9.216×10^{-7}
4	0.2	0.6	7.373×10^{-6}

Which of the following is the *correct* rate law?

- (A) $R = K[H_2] [NO]^2$
- (B) $R = K[NO]^2$

(C) $R = K[H_2]$ [NO]

(D) $R = K[H_2]^2$

72. The commutator $\left[\hat{x}, \hat{p}_{x}\right]$ is equal to :

(A) $-i\hbar$

(B) 0

(C) \hat{p}

(D) 1

73. In the Lindemann mechanism for the unimolecular reaction:

$$A(g) \ + \ M(g) \ \xleftarrow{K_1} \ A^*(g) \ + \ M(g)$$

$$A^*(g) \xrightarrow{K_2} P$$

The apparent rate constant is given by:

$$K_{obs} = \frac{K_1 K_2 [M]}{K_2 + K_{-1} [M]}$$

- (A) At high pressure the reaction becomes zeroth order w.r.t. A
- (B) At high pressure the reaction becomes first order w.r.t. M
- (C) At high pressure the reaction becomes first order w.r.t. A
- (D) Pressure has no effect on the order of reaction
- 74. The average separation of $2p_z$ electron of the H atom from the nucleus will be $(a_0$ Bohr radius):
 - (A) $3a_0/2$

(B) $5a_0$

(C) $4a_0$

- (D) a_0
- 75. Upon increasing temperature of the solution, Debye-Huckel reciprocal length (x^{-1}) will:
 - (A) Increase
 - (B) Decrease
 - (C) No change
 - (D) Initially increase, passes through maxima and then decrease

MAY - 33316/III—B

ROUGH WORK