		ooklet Code & Serial No. ाका कोड व क्रमांक <b>A</b>							
	Paper-III								
	CHEMICAL SCIENCE								
Sign	nature and Name of Invigilator	Seat No.							
1. (S	ignature)	(In figures as in Admit Ca	rd)						
(N	lame)	Seat No							
2. (Signature)		(In words)							
(N	lame)	OMR Sheet No.							
JA	N - 33318	(To be filled by the Candidate)							
	e Allowed : 2½ Hours]	[Maximum Marks : 1							
Nun	nber of Pages in this Booklet : <b>32</b>	Number of Questions in this Booklet :	75						
<ol> <li>1.</li> <li>2.</li> <li>3.</li> <li>4.</li> </ol>	<ul> <li>Instructions for the Candidates</li> <li>Write your Seat No. and OMR Sheet No. in the space provided on the top of this page.</li> <li>This paper consists of 75 objective type questions. Each question will carry twomarks. Allquestions of Paper-III will be compulsory, covering entire syllabus (including all electives, without options). At the commencement of examination, the question booklet will be given to the student. In the first 5 minutes, you are requested to open the booklet and compulsorily examine it as follows:</li> <li>(i) To have access to the Question Booklet, tear off the paper seal on the edge of this cover page. Do not accept a booklet without sticker-seal or open booklet.</li> <li>(ii) Tally the number of pages and number of questions in the booklet with the information printed on the cover page. Faulty booklets due to missing pages/ questions or questions repeated or not in serial order or any other discrepancy should not be accepted and correct booklet should be obtained from the invigilator within the period of 5 minutes. Afterwards, neither the Question Booklet will be replaced nor any extra time will be given. The same may please be noted.</li> <li>(iii) After this verification is over, the OMR Sheet Number should be entered on this Test Booklet.</li> <li>Each question has four alternative responses marked (A), (B), (C) and (D). You have to darken the circle as indicated below on the correct response against each item.</li> <li>Example : where (C) is the correct response.</li> </ul>	विद्यार्थ्यांसाठी महत्त्वाच्या सूचना 1. परिक्षार्थांनी आपला आसन क्रमांक या पृष्ठावरील वरच्या कोप-यात लि तसेच आपणांस दिलेल्या उत्तरपत्रिकेचा क्रमांक त्याखाली लिहावा. 2. सदर प्रश्नपत्रिकेत 75 बहुपर्यायी प्रश्न आहेत. प्रत्येक प्रश्नास दोन आहेत. या प्रश्नपत्रिकेती तर्मव प्रश्न सोडविणे अनिवार्य आहे. सदरचे हे या विषयाच्या संपूर्ण अभ्यासक्रमावर आधारित आहेत. 3. परीक्षा सुरू झाल्यावर विद्यार्थ्याला प्रश्नपत्रिका दिली जाईल. सुरुवाती <sup>-</sup> मिनीटांमध्ये आपण सदर प्रश्नपत्रिका उघडून खालील बाबी अवश्य तप् पहाव्यात. (i) प्रश्नपत्रिका उघडण्यासाठी प्रश्नपत्रिकवर लावलेले सील उघ सील नसलेली किंवा सील उघडलेली प्रश्नपत्रिका सिल्वघ सील नसलेली किंवा सील उघडलेली प्रश्नपत्रिकचे एकूण तसेच प्रश्नपत्रिकतील एकूण प्रश्नांची संख्या पडताळून पह पृष्ठे कमी असलेली/कमी प्रश्न असलेली/प्रश्नांचा चूव कम असलेली किंवा इतर त्रुटी असलेली सदोष प्रश्नपत्रिका ब्ल सिळणार नाही तसेच वेळही वाढवून मिळणार नाही याची वृ विद्यार्थांनी नोंद घ्यावी. (iii) वरीलप्रमाणे सर्व पडताळून पहिल्यानंतरच प्रश्नपत्रिका स्रज्यततीच्या 5 मिनिटातच पर्यवेक्षकाला परत देऊन त् प्रश्नयत्रिका मागवून घ्यावी. त्यानंतर प्रश्नपत्रिका बल मिळणार राही तसेच वेळही वाढवून मिळणार नाही याची वृ विद्यार्थांनी नोंद घ्यावी. (iii) वरीलप्रमाणे सर्व पडताळून पहिल्यानंतरच प्रश्नपत्रिक ओ.एम.आर. उत्तरपत्रिकेचा नंबर लिहावा. 4. प्रत्येक प्रश्नासाठी (A), (B), (C) आणि (D) अशी चार विकल्प उत्तरे आहेत. त्यातील योग्य उत्तर असेल तर.	न गुण च्या 5 च्या 5 पासून रा <b>पृष्ठे</b> दुसरी दुसरी दुसरी देली देली						
5.	Your responses to the items are to be indicated in the <b>OMR</b> Sheet given inside the Booklet only. If you mark at any place	A B D							
6. 7. 8. 9. 10. 11. 12.	other than in the circle in the OMR Sheet, it will not be evaluated. Read instructions given inside carefully. Rough Work is to be done at the end of this booklet. If you write your Name, Seat Number, Phone Number or put any mark on any part of the OMR Sheet, except for the space allotted for the relevant entries, which may disclose your identity, or use abusive language or employ any other unfair means, you will render yourself liable to disqualification. You have to return original OMR Sheet to the invigilator at the end of the examination compulsorily and must not carry it with you outside the Examination Hall. You are, however, allowed to carry the Test Booklet and duplicate copy of OMR Sheet on conclusion of examination. <b>Use only Blue/Black Ball point pen.</b> <b>Use of any calculator or log table, etc., is prohibited.</b> <b>There is no negative marking for incorrect answers.</b>	<ol> <li>या प्रश्नपत्रिकेतील प्रश्नांची उत्तरे ओ.एम.आर. उत्तरपत्रिकेतच दर्शवा इतर ठिकाणी लिहीलेली उत्तरे तपासली जाणार नाहीत.</li> <li>आत दिलेल्या सूचना काळजीपूर्वक वाचाव्यात.</li> <li>प्रश्नपत्रिकेच्या शेवटी जोडलेल्या कोन्या पानावरच कच्चे काम करावे.</li> <li>जर आपण ओ.एम.आर. वर नमूद केलेल्या ठिकाणा व्यतिरीक्त इतर क नाव, आसन क्रमांक, फोन नंबर किंवा ओळख पटेल अशी कोणतीही केलेली आढळून आल्यास अथवा असभ्य भाषेचा वापर किंवा इतर गैरमा अवलंब केल्यास विद्यार्थ्याला परीक्षेस अपात्र ठरविण्यात येईल.</li> <li>परीक्षा संपल्यानंतर विद्यार्थ्याने मूळ ओ.एम.आर. उत्तरपत्रिका पर्यवेक्षक परत करणे आवश्यक आहे. तथापी, प्रश्नपत्रिका व ओ.एम.आ. उत्तरपत्रि द्वितीय प्रत आपल्याबरोबर नेण्यास विद्यार्थ्यांना परवानगी आहे.</li> <li>फक्त निळ्या किंवा काळ्या बॉल पेनचाच वापर करावा.</li> <li>कॅलबयुलेटर किंवा लॉग टेबल वापरण्यास परवानगी नाही.</li> <li>चुकीच्या उत्तरासाठी गुण कपात केली जाणार नाही.</li> </ol>	1. कोठेही ो खूण गार्गांचा कांकडे						

# Chemical Science Paper III

# Time Allowed : 2½ Hours][Maximum Marks : 150Note : This paper contains seventy five (75) multiple choice questions, each<br/>question carrying Two (2) marks. Attempt All questions.

- 1. The *correct* statement for atomic absorption spectroscopy (AAS) from the following :
  - (A) Non-metals are determined by direct methods
  - (B) Beer's law is not obeyed over wide concentration range
  - (C) Separate lamp for each element to be determined is required
  - (D) Al, Ti, W, Mo and Si can be detected by AAS
- 2. The capacity to undergo exchange reaction by the hydrated metal ions of the same size in ion exchange chromatography follows the order :

(A) 
$$Al^{3+} > Ca^{2+} > Th^{4+} > Na^{+}$$

(B) 
$$Th^{4+} > Al^{3+} > Ca^{2+} > Na^+$$

(C) 
$$Na^+ > Ca^{2+} > Al^{3+} > Th^{4+}$$

(D) 
$$Ca^{2+} > Al^{3+} > Th^{4+} > Na^{+}$$

3. The thermal methods (I) and their parameters (II) are given below :

I		II	
( <i>a</i> ) TG	( <i>e</i> )	$\Delta T$	
(b) DTG	( <i>f</i> )	dm/dt	
(c) DTA	( <i>g</i> )	dH/dt	
(d) DSC	( <i>h</i> )	Mass	
The <i>correct</i> match is :			
(A) $(a)$ - $(e)$ , $(b)$ - $(f)$ , $(c)$ - $(g)$ , $(d)$ - $(h)$			
(B) $(a)-(f), (b)-(g), (c)-(h), (d)-(e)$			
(C) $(a)-(h), (b)-(f), (c)-(e), (d)-(g)$			
(D) $(a)-(g), (b)-(e), (c)-(f), (d)-(h)$			

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- The intervalence band in Mixed-valent compounds usually occurs in the ..... 4. region.
  - (A) Ultra-violet (B) Visible
  - (C) Near infrared (D) Far infrared

In the base-catalysed substitution of  $\rm Cl^-$  by  $\rm OH^-$  in  $\rm [Co(\rm NH_3)_5 Cl]^{2+}$  under 5. strongly basic conditions, the first step in the mechanism is :

- (A) Conversion of an ammine to amido ligand
- (B) Substitution of Cl<sup>-</sup> by OH<sup>-</sup>
- (C) Dissociation of Cl<sup>-</sup> to give 5-coordinate intermediate
- (D) Assocaition of OH<sup>-</sup> to give 7 coordinate intermediate

To satisfy the 18 electron rule the value of *n* in the complex  $[Rh(\eta^5 - C_5h_5)(CO)_n]$ 6. should be :

- (A) 2 (B) 3
- (C) 1 (D) 4
- 7. The correct order of Nephelauxetic parameter  $(\beta)$  of the ligands is :

(A) 
$$CN^{-} > NC\overline{S} > en > NH_3 > H_2O$$
  
(B)  $H_2O > NH_3 > en > NC\overline{S} > CN^{-}$ 

- $(C) \quad en > CN^- > NCS^- > NH_3 > H_2O$
- $(D) \quad H_2O > NH_3 > NCS^- > CN^- > en$
- Jahn-Teller distortion will be exhibited by : 8.
  - (A)  $[Mn(H_2O)_6]^{2+}$ (B)  $[Cr(H_2O)_6]^{3+}$
  - $(C) \ [Ni(H_2O)_6]^{2+}$  $(D) \quad [Cu(H_2O)_6]^{2+}$

- 9. Rieske protein is a [2Fe-2S] cluster wherein :
  - (A) One Fe atom is coordinated to two terminal imidazole ligands and the other Fe atom is coordinated to two terminal cystine ligands
  - (B) Both Fe atoms are coordianted to two terminal imidazole groups each
  - (C) Both Fe atoms are coordianted to two terminal cystine groups each
  - (D) Both Fe atoms are coordianted to one terminal imidazole and one cystine each

10.  $[Ni(dipyridyl)_3]SO_4$  records three transitions in electronic spectra at 12,650 cm<sup>-1</sup>, 19,200 cm<sup>-1</sup> and 26,000 cm<sup>-1</sup>. The 10 Dq of the complex is :

- (A)  $12650 \text{ cm}^{-1}$  (B)  $15180 \text{ cm}^{-1}$
- (C) 10120  $\rm cm^{-1}$  (D) 26000  $\rm cm^{-1}$
- 11. In which of the following organometallic complexes, the carbon based ligands is both  $\sigma$  and  $\pi$  bonded.

12. The EPR spectra of  $[Mn(H_2O)_6]^{2+}$  (<sup>55</sup>Mn=I=5/2) with zero field splitting show five set of lines. Each set further consist of ...... .... hyperfine lines (At. No. Mn = 25)

- (A) 5 (B) 6
- (C) 30 (D) 1

13. In the Wacker catalystic cycle if methanol is used in place of water as nucleophile the product will be :

- (A) Ethanol (B) Methyl vinyl ether
- (C) Acetaldehyde (D) Acetone

14. 'A' type of ground Mülliken symbol in complex having oh symmetry is :

		0	
	sphere electron transfer reaction ?		
	(A) $[Co(NH_3)_5NO_3]^{2+}/[V(OH_2)_6]^{2+}$		
	(B) $[C_0(NH_3)_5I]^{2+}/[V(OH_2)_6]^{2+}$		
	(C) $[Co(en)_3]^{3+}/[V(OH_2)_6]^{2+}$		
	(D) $[Co(NH_3)_5Br]^{2+}/[V(OH_2)_6]^{2+}$		
16.	16. The $d^n$ electron configuration of the first row transition	metal ions that show	
	'high-spin'—'low-spin' transition are :		
	(A) $d^4$ and $d^5$ (B) $d^5$ and $d^6$		
	(C) $d^4$ , $d^5$ , $d^6$ and $d^7$ (D) $d^4$ , $d^5$ and	$d^6$	
17.	17. The measurement of the intensity of the scattered ligh	nt as a function of the	
	concentration of the dispersed phase form is the princ	piple of :	
	(A) Colorimetry (B) Turbidimet	ry	
	(C) Nephelometry (D) Phosphorin	netry	
18.	18. Racemization of octahedral complexes containing three	bidentate rings by the	
	Ray-Dutt twist mechanism forms intermediates of	point group.	
	(A) $D_{3h}$ (B) $C_{2v}$		
	(C) $C_{3v}$ (D) $D_{2h}$		
19.	19. The $^{31}P{OH}$ NMR spectra of $HPO(OH)_2$ and $H_2PO(OH)$	$(I = {}^{3}p, {}^{1}H = 1/2)$ shows	
	respectively :		
	(A) doublet and triplet (B) triplet and	triplet	
	(C) triplet and doublet (D) doublet and	d doublet	
20. The following statement is <i>incorrect</i> with respect to ${}^{57}$ Fe Mössb		Fe Mössbauer spectra	
	of $FeCl_3.6H_2O$ and $FeCl_2.4H_2O$ :		
	(A) Both the compounds show doublet with equal qua	adrupole splitting	
	(B) Both are compounds show doublet with unequal quadrupole splitting		
	(C) Both the compounds are high spin complexes of $\ensuremath{Fe}(II)$ and $\ensuremath{Fe}(III)$		
	respectively		
	(D) Valence contribution is present in $FeCl_2.4H_2O$ wh	ile lattice contribution	
	is present in $\text{FeCl}_3.6\text{H}_2\text{O}$		
	6		

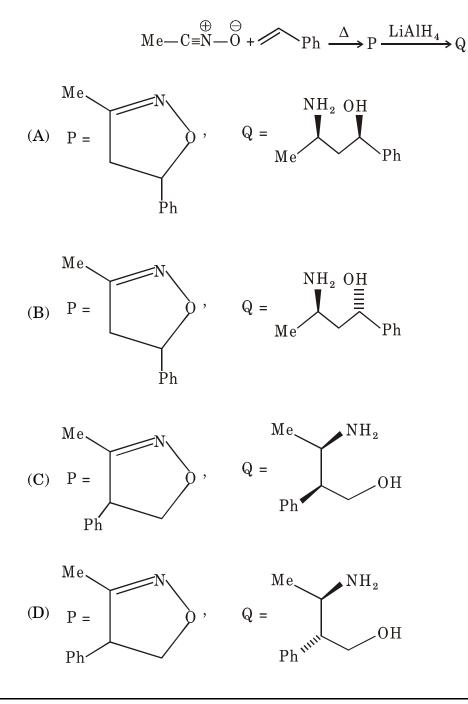
15. Which of the following pairs of coordination complexes do not undergo inner

- 21. Which of the following pairs of elements have  $(n-2)f^7(n-1)d^1ns^2$  electronic configuration ?
  - (A)  $S_m$ ,  $C_m$  (B)  $G_d$ ,  $C_m$
  - (C)  $\mathbf{E}_{\mathbf{u}}, \mathbf{C}_{\mathbf{m}}$  (D)  $\mathbf{E}_{\mathbf{r}}, \mathbf{C}_{\mathbf{m}}$
- 22. Which of the following statements with respect to 'trans effect' is true ?
  - (A) Ammonia > Chloride (B) Water > Chloride
  - (C) Pyridine > Chloride (D) Chloride > Ammonia
- 23. In the layered structure of  $CdI_2$ ,  $Cd^{2+}$  occupies ..... octahedral sites.

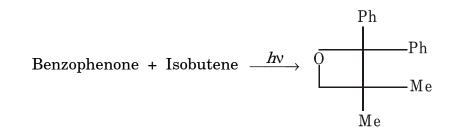
(A) 
$$\frac{1}{2}$$
 (B)  $\frac{1}{3}$   
(C)  $\frac{1}{4}$  (D) 1

- 24. A dimeric Ruthenium (IV) complex [K<sub>4</sub>(Cl<sub>5</sub>Ru—O—RuCl<sub>5</sub>] is diamagnetic at room temperature (26°C) althrough there are two unpaired electrons per Ru(IV) ion, the magnetic behaviour can be best explained by :
  - (A) Antiferromagnetic coupling (B) Ferromagnetic coupling
  - (C) Ferrimagnetic coupling (D) Antiferrimagnetic coupling
- 25. Tetragonal distorted complexes will be exhibited by :
  - (A) Low spin Fe<sup>2+</sup> (B) High spin Cr<sup>2+</sup>
  - (C) High spin  $Mn^{2+}$  (D)  $Ni^{2+}$

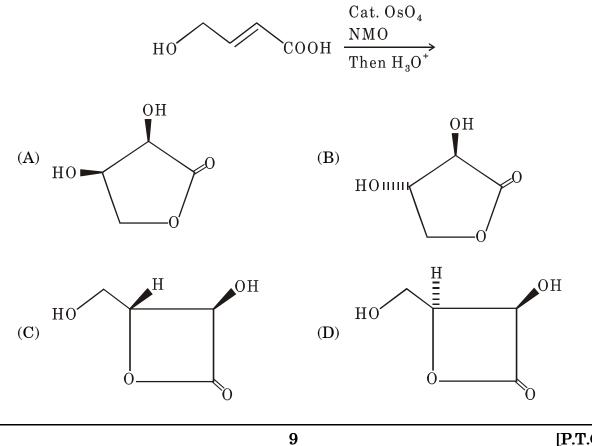
26. The major products P and Q formed in the following reaction sequence are :



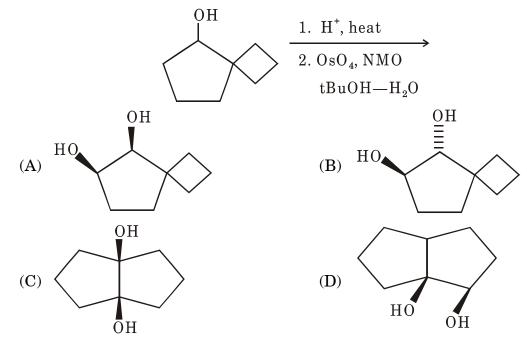
27. The following reaction is an example of :



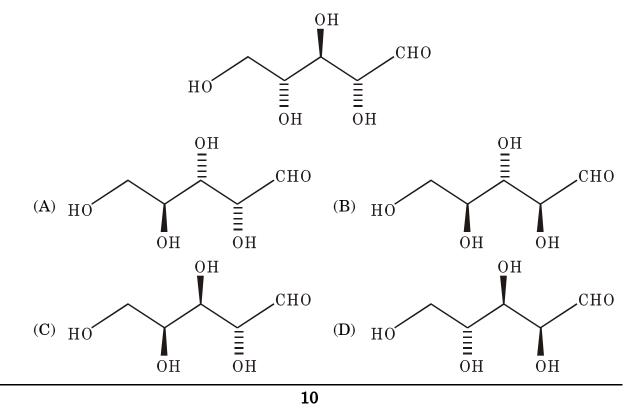
- (A) Norrish type I reaction (B) Norrish type II reaction
- (C) Borton reaction (D) Paterno-Buchi reaction
- 28. The major product formed in the following reaction is :

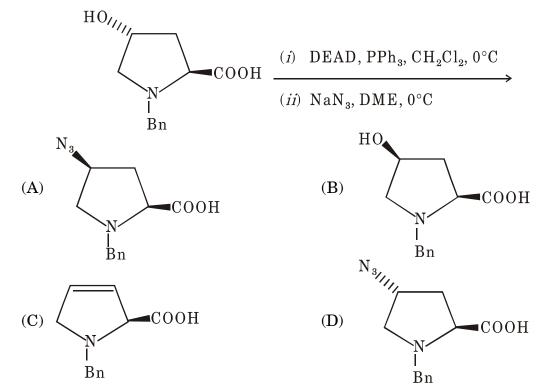


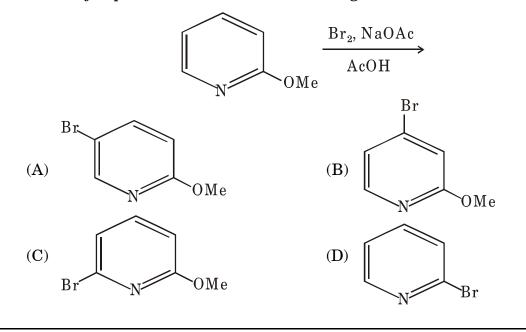
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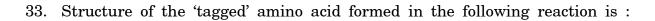


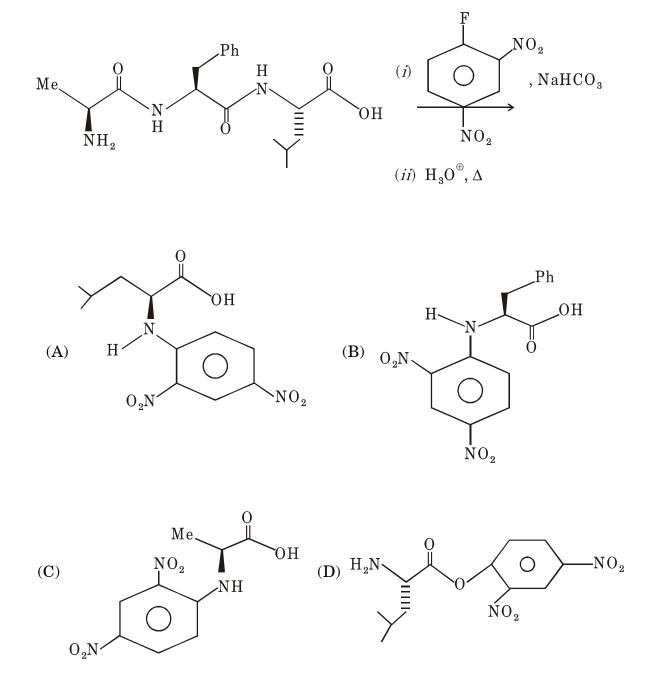
30. Find out the correct structure of L-Ribose if the structure of D-Ribose is :



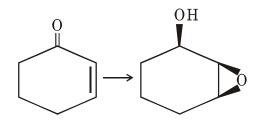




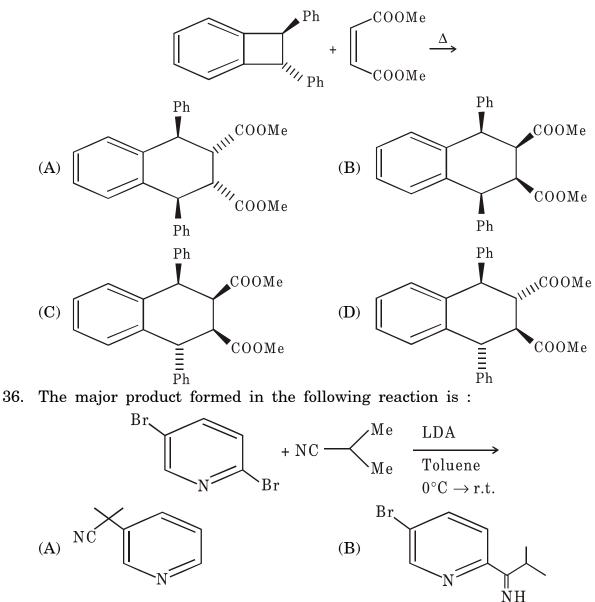


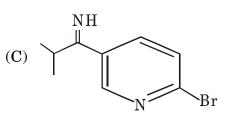


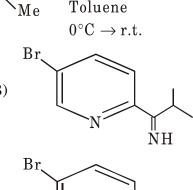
34. The correct reagents to effect the following conversion are :

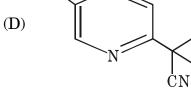


- (A) (i)  $NaBH_4$ , CeCl<sub>3</sub>, MeOH
  - (*ii*) m—CPBA,  $CH_2Cl_2$ , 0°C
- (B) (i) NaBH<sub>4</sub>, CeCl<sub>3</sub>, MeOH
  - (ii) H<sub>2</sub>O<sub>2</sub>, NaOH, 0°C
- (C) (i) m—CPBA,  $CH_2Cl_2$ , 0°C
  - (*ii*) LiAlH<sub>4</sub>, THF,  $0^{\circ}C \rightarrow rt$
- (D) (i)  $H_2O_2$ , NaOH, 0°C  $\rightarrow$  rt
  - (*ii*) LiAlH<sub>4</sub>, THF,  $0^{\circ}C \rightarrow \text{reflux}$

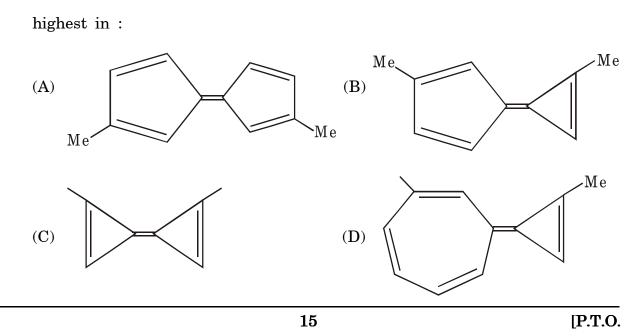








14



37. The correct match of natural products in Column-I with class in Column-II is :

#### Column-I

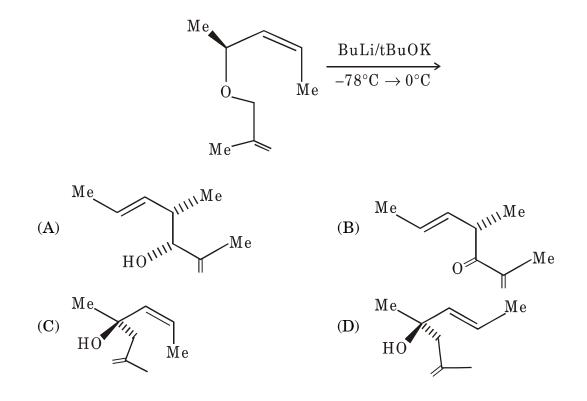
(P) Cholesterol

- Column-II
- (i) Polysaccharide
- (Q) Cellulose (*ii*) Steroid
- $(\mathbf{R})$  Quinine

(iii) Vitamin

- (*iv*) Alkaloid
- (A) (P)-(*iii*), (Q)-(*i*), (R)-(*iv*)
- (B) (P)-(ii), (Q)-(iv), (R)-(i)
- (C) (P)-(*ii*), (Q)-(*i*), (R)-(*iv*)
- (D) (P)-(*i*), (Q)-(*iii*), (R)-(*iv*)

38. Amongst the following the rate of cis-trans isomerisation is expected to be

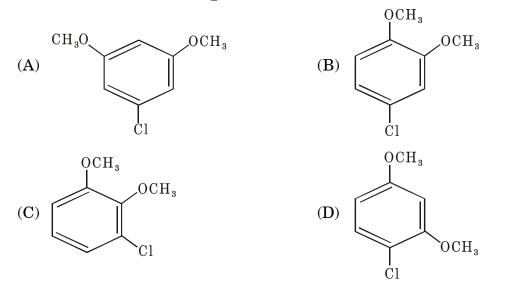


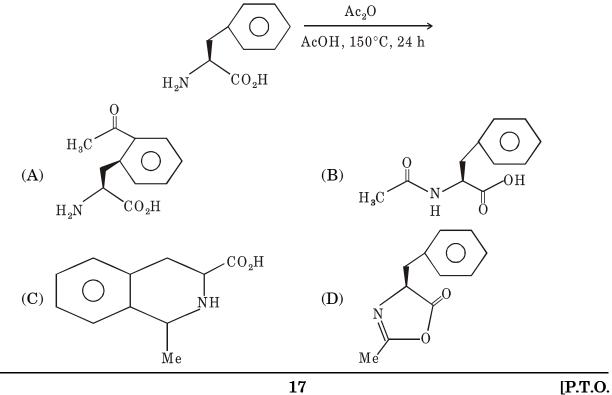
40. The proton decoupled  $^{13}C$  NMR spectrum of  $HF_2C$ — $CH_2Cl$  will show :

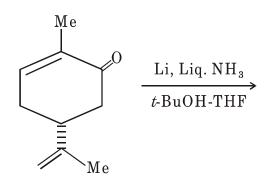
- (A)  ${\rm C}_1$  as singlet and  ${\rm C}_2$  as singlet
- (B)  $C_1$  as triplet and  $C_2$  as triplet
- $(C)\ C_1$  as triplet and  $C_2$  as singlet
- (D)  ${\rm C}_1$  as doublet and  ${\rm C}_2$  as singlet

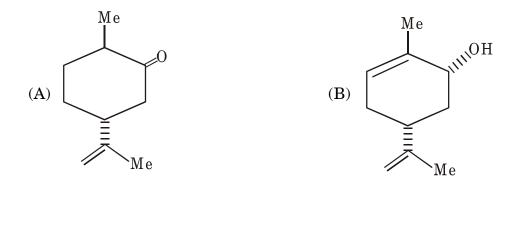
The correct structure of a compound that exhibits the following <sup>1</sup>H-NMR spectral 41. data is :

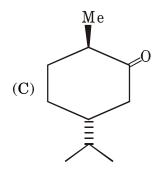
 $\delta 3.8({\rm S},\, 3{\rm H}),\, 3.9~({\rm S},\, 3{\rm H})$  6.82  $(dd,\, {\rm J}$  = 8 and 2  ${\rm H}_2,\, 1{\rm H}),\, 7.06~(t,\, {\rm J}$  = 8  ${\rm H}_2,\, 1{\rm H}),\, 7.15~(dd,\, {\rm J}$  = 8 and 2  ${\rm H}_2,\, 1{\rm H})$ 

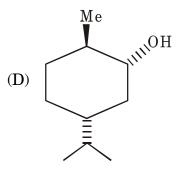


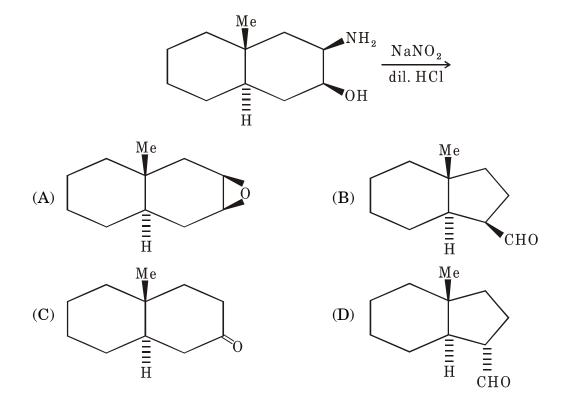




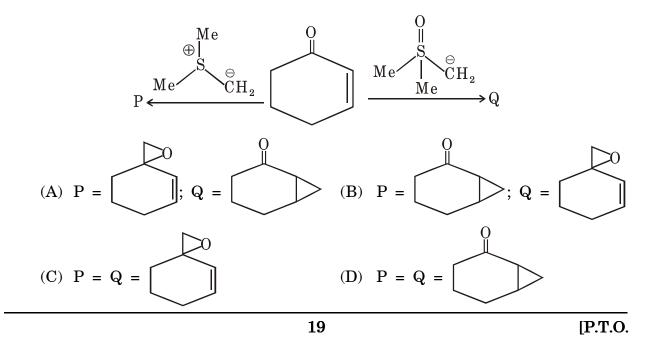


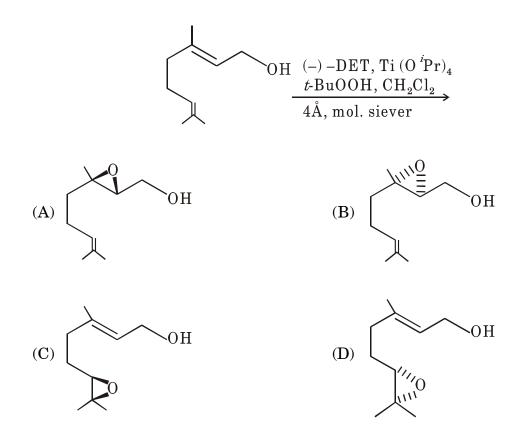


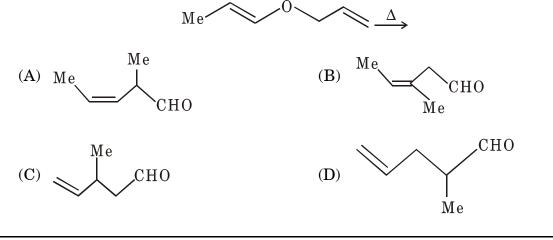


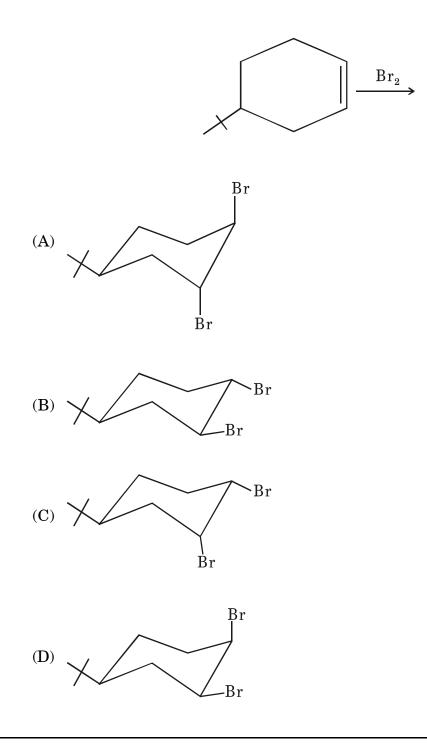


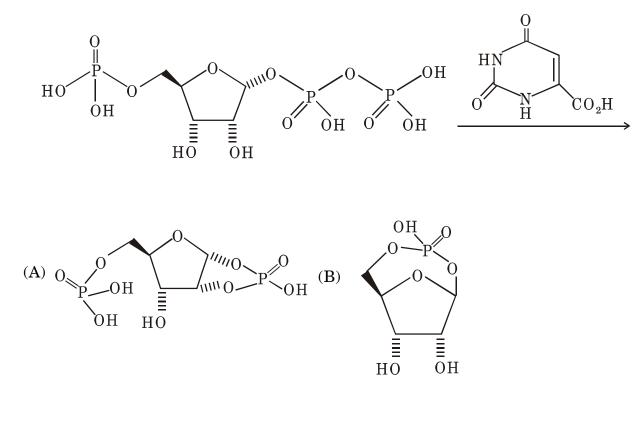
45. Major product formed in the following reaction is :

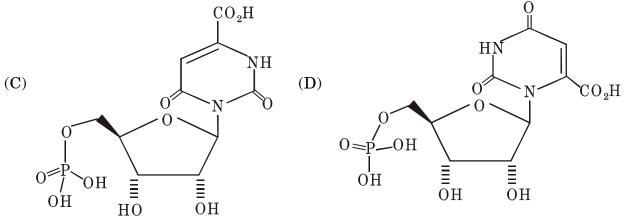




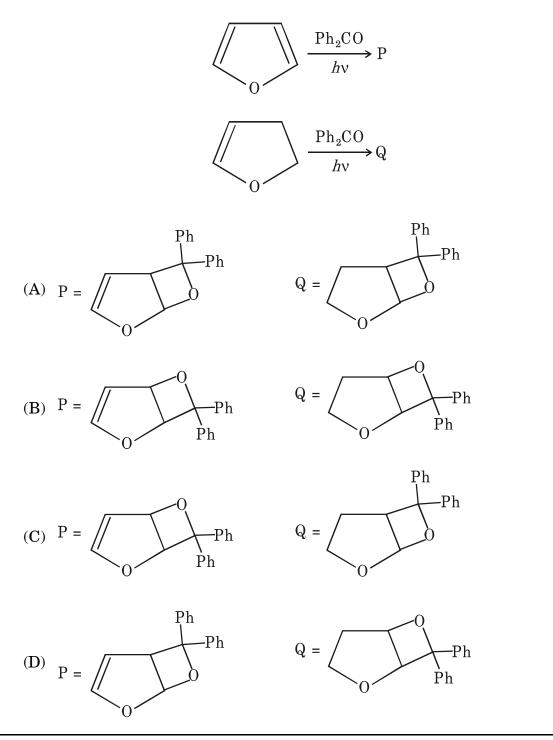








50. The major product P and Q formed in the following reactions are :



51. One of the most widely used functions for the van der Waals potential is the Lennard-Jones expressed as :

(A) 
$$\mathbf{E}_{\mathrm{LJ}} = \sum_{\substack{\text{nonbonded} \\ \text{pairs}}} \left( \frac{\mathbf{A}_{ik}}{\mathbf{r}_{ik}^{6}} - \frac{\mathbf{C}_{ik}}{\mathbf{r}_{ik}^{12}} \right)$$
(B) 
$$\mathbf{E}_{\mathrm{LJ}} = \sum_{\substack{\text{nonbonded} \\ \text{pairs}}} \left( \frac{\mathbf{A}_{ik}}{\mathbf{r}_{ik}^{12}} - \frac{\mathbf{C}_{ik}}{\mathbf{r}_{ik}^{6}} \right)$$
(C) 
$$\mathbf{E}_{\mathrm{LJ}} = \sum_{\substack{\text{nonbonded} \\ \text{pairs}}} \left( \frac{\mathbf{A}_{ik}}{\mathbf{r}_{ik}^{12}} + \frac{\mathbf{C}_{ik}}{\mathbf{r}_{ik}^{6}} \right)$$
(D) 
$$\mathbf{E}_{\mathrm{LJ}} = \sum_{\substack{\text{nonbonded} \\ \text{pairs}}} \left( \frac{\mathbf{A}_{ik}}{\mathbf{r}_{ik}^{6}} + \frac{\mathbf{C}_{ik}}{\mathbf{r}_{ik}^{12}} \right)$$

52. Coefficient of thermal expansion ( $\alpha$ ) is given by :

53. The distribution function for identical indistinguishable particles can be

represented by the equation  $N_j = \frac{g_j}{e^{(\in_j - \mu)/kT} + a}$ , which of the following statements

is correct?

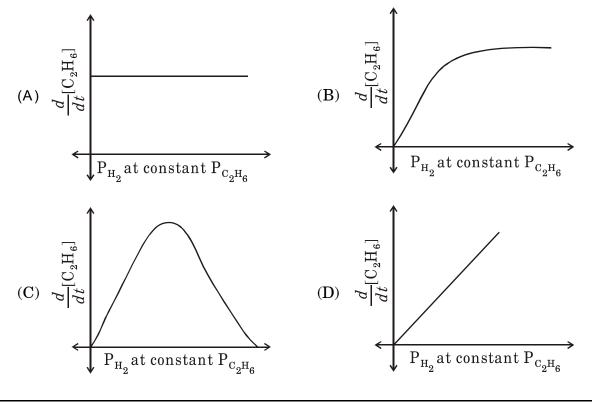
- (A) a = +1 for Bose-Einstein statistics
- (B) a = -1 for Fermi-Dirac statistics
- (C) a = +1 for Fermi-Dirac statistics
- (D) a = 1/2 for Bose-Einstein statistics

#### 54. For an isolated system :

- (A)  $d\mathbf{S} = d_e \mathbf{S}$  (B)  $d\mathbf{S} = d_I \mathbf{S}$
- (C)  $d_e S > 0$  (D)  $d_e S < 0$

- 55. If there is no net flow of electrical current, then generation of potential difference per unit change of pressure and generation of potential difference per unit change of temperature is termed as ......... respectively.
  - (A) Thermoelectric power and osmotic pressure
  - (B) Osmotic pressure and thermoelectric power
  - (C) Streaming potential and thermoelectric power
  - (D) Thermoelectric power and streaming potential
- 56. For any polydispersed system, which of the following statements is true?
  - (A) Number average relative molar mass is greater than mass average relative molar mass
  - (B) Number average relative molar mass is less than mass average relative molar mass
  - (C) Number average relative molar mass coincide with mass average relative molar mass
  - $(D) \ None \ of \ the \ above$
- 57. A particle in a one-dimensional box of length L can be excited from n = 1 to n = 2 state with frequency v. If the box length is doubled, the frequency needed to achieve this transition is :
  - (A)  $\nu/4$  (B)  $\nu/2$
  - (C) 2v (D) 4v
- 58. If operator  $\hat{D}^n(n \text{ times differentiate with respect to } x)$  operates on eigenfunction  $e^{ax}$ , the eigenvalue would be :
  - (A)  $n\alpha$  (B)  $n^{\alpha}$
  - (C)  $\alpha^n$  (D)  $n!\alpha$
- 59. The eigenfunctions of a rigid rotator are :
  - (A) Spherical harmonics (B) Laguerre polynomials
  - (C) Hermite polynomials (D) Legendre transformers

- 60. According to Hund's rule, most stable state for  $np^2$  configuration is .....
  - (A)  ${}^{1}S_{0}$  (B)  ${}^{3}P_{0}$
  - (C)  ${}^{3}P_{1}$  (D)  ${}^{3}P_{2}$
- 61. According to simple HMO theory, which one of the following statements about butadiene is *true* ?
  - (A) It has a non-bonding MO
  - (B) Exciting an electron from the HOMO to the LUMO will not change the  $\pi$  bond orders
  - (C) Exciting an electron from the HOMO to the LUMO will not change the  $\pi$  charge densities
  - (D) It is most likely to undergo electrophilic substitution at one of the two inner carbons
- 62. In the hydrogenation of  $C_2H_4$  on a Ni surface both the alkene and  $H_2$  are adsorbed on similar active sites of the catalyst surface. Which of the following figures depict the reaction ?

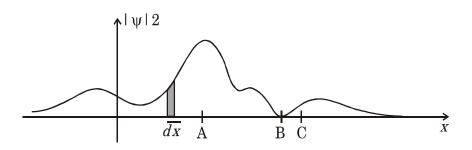


- 63. In case of heterogeneous catalysis which of the following is correct?
  - (A) At least one reactant must be physisorbed on the catalyst
  - (B) At least one reactant must be chemisorbed on the catalyst
  - (C) The number of active sites decrease with increase in surface area
  - (D) Surface area can be determined by spectroscopic studies
- 64. A surface of Li metal (sample X) was suspected to be completely coated with a thin layer of Li<sub>2</sub>O (sample Y). When X and Y are analysed by X-ray photoelectron spectroscopy :
  - (A) Kinetic energy of the photoelectrons from X will be higher than that from Y
  - (B) Kinetic energy of the photoelectrons from Y will be higher than that from Y
  - (C) Binding energy of the photoelectrons from Y will be lower than that from X
  - (D) Kinetic energy of the photoelectrons from both X and Y will be the same
- 65. A reactor vessel at 1000 K contains  $CO_2$  at a pressure of 0.5 atm. Part of the  $CO_2$  is converted into CO by the addition of graphite. If the total pressure in the reactor vessel at equilibrium is 0.8 atm, the value of  $K_P$  for the reaction of  $CO_2$  with graphite is :
  - (A) 1.8 (B) 3.24
  - (C) 0.324 (D) 09

66. For the equilibrium  $Fe(s) + H_2O(g) \rightleftharpoons FeO(s) + H_2(g)$ , the number of components, phases and degrees of freedom are respectively :

(A) 3, 3, 2	(B) 3, 4, 2
(C) 3, 3, 3	(D) 3, 2, 1

## 67. Given below is the wave function of a particle :



Which of the following is *correct*?

- (A) The particle is more likely to be found near B
- (B) The particle is more likely to be found near A
- (C) The shaded area represents the energy of the particle when it travels through a distance dx
- (D) The shaded area represents the amplitude of the particle in an area dx
- 68. The de Broglie wavelength of an electron travelling with 1% speed of light is about :

(mass of electron and speed of light are 9.109  $\times$   $10^{-31}$  kg and 2.998  $\times$   $10^8~ms^{-1}$  respectively) :

- (A) 243 m (B) 243 cm
- (C) 243 Å (D) 243 pm

69. Surface tension of  $Br_2(I)$  water and mercury are 41.0 mNm<sup>-1</sup>, 72.0 mN/m and 486.2 mNm<sup>-1</sup> respectively at 25°C. These values imply that :

- (i) Metallic bonding is weaker than covalent bonding
- (ii) Hydrogen bonding is stronger than van der Waals bonding
- (iii) Metallic bonding is stronger than hydrogen bonding
- (iv) All types of intermolecular forces are of similar strength
- (A) (iii) and (iv) (B) (ii) and (iii)
- (C) (i), (ii) and (iii) (D) (iv), (ii) only

70. In case of radical polymerization, the kinetic chain length is given by expression :

$$\nu = \frac{k_p [\mathbf{M}] [\mathbf{I}]^{-1/2}}{2k_t f^{1/2} \left(\frac{k_d}{k_t}\right)^{1/2}}$$

increase in the initiator concentration eventually leads to :

- (A) Higher molecular weight polymer
- (B) Low molecular weight polymer
- (C) Rate of reaction and thus the extent polymerization does not depend on the concentration of initiator
- (D) Fast polymerization reaction
- 71. Consider the following reaction in an aqueous solution :

 $I^{-}(aq) + OCl^{-}(aq) \rightarrow IO^{-}(aq) + Cl^{-}(aq)$ 

and the following initial concentration and initial rate data for this reaction :

[I <sup>-1</sup> ], M	[OCl <sup>-</sup> ], M	Initial Rate, $Ms^{-1}$
0.1	0.05	$3.05*10^{-4}$
0.2	0.05	$6.10*10^{-4}$
0.3	0.01	$1.83 * 10^{-4}$
0.4	0.02	$6.12*10^{-4}$

Which of the following is the *correct* rate law for this reaction ?

(A) Rate =  $k[I^{-}]^{2}[OCI^{-}]$ 

- (B) Rate =  $k[OCl^{-}]$
- (C) Rate =  $k[I^{-}]^{2}$
- $(\mathbf{D}) \quad \mathbf{M} = \mathbf{M} \begin{bmatrix} \mathbf{0} \\ \mathbf{0} \end{bmatrix}$
- (D) Rate =  $k[I^-][OCI^-]$

[P.T.O.

72. Consider the following electrochemical cell, connected to the potential source (Potentiostat) with Hg as a working electrode :

when the potential is change towards (i) more negative direction and (ii) change towards the more positive direction from the equilibrium potential value, the following observations are expected :

Given :

$$\begin{split} & E^{\circ}(Cd^{2+}/Cd^{\circ}) \; = \; - \; 0.351 \ V, \\ & E^{\circ}(Zn^{2+}/Zn^{\circ}) \; = \; - \; 0.762 \ V, \; E^{\circ}(Hg^{2+}/Hg^{\circ}) \; = \; + \; 0.855 \ V \end{split}$$

- (A) (i)  $H_2$  evolution will start.
  - (ii) O<sub>2</sub> evolution will start.
- (B) (i) Cd will deposit first followed by Zn.
  - (*ii*) Hg will start oxidizing.
- (C) (*i*) Zn will deposit.
  - (ii) O<sub>2</sub> evolution will be observed.
- (D) (*i*) Cd will deposit.
  - (*ii*) Hg will start oxidizing.

- 73. The ideal solution made up of 5.00 mol (mole fraction = 0.606) benzene and 3.25 mol (mole fraction = 0.394) toluene, the free energy of mixing ( $\Delta G_{mix}$ ) is equal to -13 kJ mol<sup>-1</sup>. The corresponding values for the their equimolar solution will be ;
  - (A) Unity
  - (B)  $(\Delta G_{mix})$  does not depend on mole fraction, thus equal to  $-13~kJ~mol^{-1}$
  - (C) Less negative than that of  $-13\ kJ\ mol^{-1}$
  - (D) More negative than that of  $-13 \text{ kJ mol}^{-1}$

74. The ideal solution made up of 5.00 mol (mole fraction = 0.606) benzene and 3.25 mol (mole fraction = 0.394) toluene, the change in enthalpy during mixing  $(\Delta H_{mix})$  would be :

- (A) zero (B)  $1.0 \text{ kJ mol}^{-1}$
- (C) positive (D) negative
- 75. An electrochemical cell is constructed using half cell for the reduction reaction is given by :

 $Fe(OH_2)(s)$  + 2e  $\rightarrow$  Fe(s) + 2OH^-(aq); E^{\circ} = –0.877 V

It is combined with another half cell for which the cell reaction is given by :

$$Al^{3+}(aq) + 3e \rightarrow Al(s); E^{\circ} = -1.66 V$$

Activities of all the components are assumed to be unity.

The overall reaction for the cell in the direction of spontaneous change is :

$$\begin{array}{ll} (A) & (Fe(OH)_2(s) \,+\, Al(s) \rightarrow Fe(s) \,+\, OH^-(aq.) \,+\, Al^{3+} \,\, (aq.) \\ (B) & 2(Fe(OH)_2(s) \,+\, 3Al(s) \rightarrow 2Fe(s) \,+\, 6OH^-(aq.) \,+\, 3Al^{3+} \,\, (aq.) \\ (C) & 3(Fe(OH)_2(s) \,+\, 2Al(s) \rightarrow 3Fe(s) \,+\, 6OH^-(aq.) \,+\, 2Al^{3+} \,\, (aq.) \\ (D) & Fe(s) \,+\, OH^-(aq.) \,+\, Al^{3+}(aq.) \rightarrow (Fe(OH)_2(s) \,+\, Al(s) \\ \end{array}$$

**ROUGH WORK**