



INDIAN SCHOOL AL WADI AL KABIR

REHEARSAL EXAMINATION-I

08/12/2022

CHEMISTRY (043)

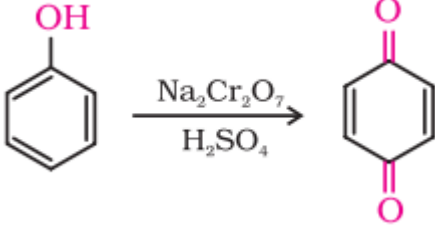
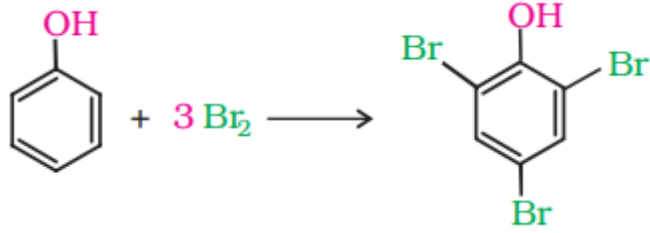
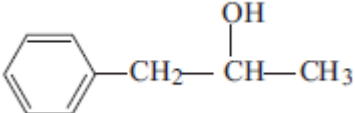
Maximum Marks: 70

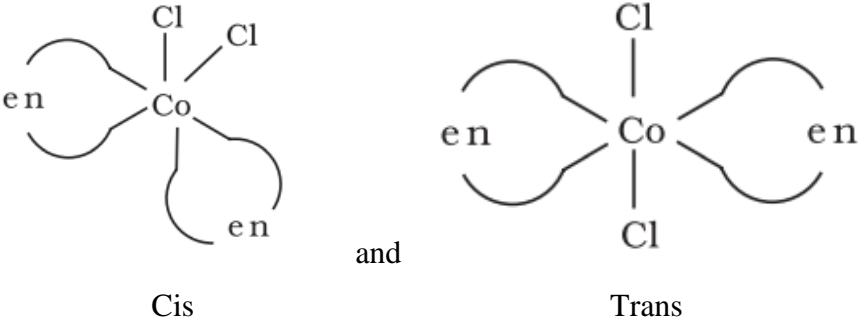
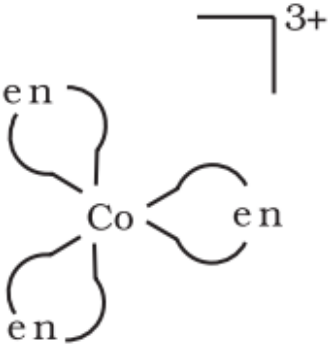
Class: XII

Time: 3 Hours

SET – I – ANSWER KEY

1.	(A) osmotic pressure	1
2.	(C) first order reaction	1
3.	(B) Argon	1
4.	(C) (iii) and (iv)	1
5.	(D) 2	1
6.	(D) +3	1
7.	(A) i, iv	1
8.	(D) Benzyl halides are more reactive than vinyl and aryl halides	1
9.	(B) a dehydrohalogenation reaction	1
10.	(B) Scandium	1
11.	(C) Linkage Isomers	1
12.	(C) Polypeptides	1
13.	(A) Acetone	1
14.	(B) 2-Methylbutan-2-ol	1
15.	(C) Assertion is correct statement but reason is wrong statement.	1
16.	(D) Assertion is wrong statement but reason is correct statement.	1
17.	(A) Assertion and reason both are correct statements and reason is correct explanation for assertion.	1
18.	(C) Assertion is correct statement but reason is wrong statement.	1
19.	a) Oxygen stabilizes Mn more than F due to multiple bonding.	1
	b) This is due to decrease in size and increase in mass from titanium to copper.	1
20.	$k = 0.693/t$	$\frac{1}{2}$
	$k = 0.0277 \text{ min}^{-1}$	$\frac{1}{2}$
	$t_{80\%} = (2.303/0.0277) \log 100/20$	$\frac{1}{2}$
	$= 58.11 \text{ min}$	$\frac{1}{2}$

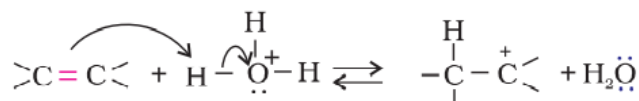
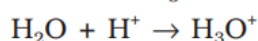
21.	<p>a) In phenol, lone pair of electrons on oxygen are delocalized over benzene ring due to resonance but in alcohol lone pair of electrons on oxygen are localized and hence available for protonation.</p> <p>b) In anisole, O-C₆H₅ bond is stronger than O-CH₃ bond as O-C₆H₅ bond has partial double bond character due to resonance.</p>	1 1
22.	<p>a) </p> <p>b) </p> <p style="text-align: center;">OR</p> <p>a) i) PCC (or any other suitable reagent) ii) Conc. HNO₃</p> <p>b) </p>	1 1 1/2 1/2 1
23.	<p>Due to osmosis.</p> <p>An increase in temperature would accelerate the process of osmosis.</p>	1 1
24.	<p>a) [Co(NH₃)₄(H₂O)Cl]Cl₂</p> <p>b) Coordination number is 6 Oxidation state of chromium is +3</p> <p style="text-align: center;">OR</p>	1 1/2 1/2

	<p>a) </p> <p style="text-align: center;">Cis and Trans</p> <p>b) </p>	<p style="text-align: right;">$\frac{1}{2} + \frac{1}{2}$</p> <p style="text-align: center;">1</p>
25.	<p>a) 2,3-Dimethylbutane</p> <p>b) Isopropyl chloride < 1-Chloropropane < 1-Chlorobutane</p>	<p style="text-align: right;">1</p> <p style="text-align: right;">1</p>
26.	<p>a) Order of reaction with respect to A is 1</p> <p style="padding-left: 2em;">Order of reaction with respect to B is 0</p> <p>b) Rate law is, Rate = k[A]</p> <p style="padding-left: 2em;">Overall order of reaction is 1</p> <p>c) II</p>	<p style="text-align: right;">$\frac{1}{2}$</p> <p style="text-align: right;">$\frac{1}{2}$</p> <p style="text-align: right;">$\frac{1}{2}$</p> <p style="text-align: right;">$\frac{1}{2}$</p> <p style="text-align: right;">1</p>
27.	<p>a) On addition of barium chloride, $[\text{Co}(\text{NH}_3)_5\text{Cl}]\text{SO}_4$ forms white precipitate of BaSO_4 while $[\text{Co}(\text{NH}_3)_5(\text{SO}_4)]\text{Cl}$ does not.</p> <p>b) Diamminechloridonitrito-N-platinum(II)</p> <p>c) i) Strong field ligand: $t_2g^5 e_g^0$</p> <p style="padding-left: 2em;">ii) Weak field ligand: $t_2g^3 e_g^2$</p> <p style="text-align: center;">OR</p> <p>a) Type of hybridization – sp^3d^2</p> <p>b) Magnetic property - Paramagnetic</p> <p>c) Type of complex – Outer orbital complex</p>	<p style="text-align: right;">1</p> <p style="text-align: right;">1</p> <p style="text-align: right;">$\frac{1}{2}$</p> <p style="text-align: right;">$\frac{1}{2}$</p> <p style="text-align: right;">1</p> <p style="text-align: right;">1</p> <p style="text-align: right;">1</p>

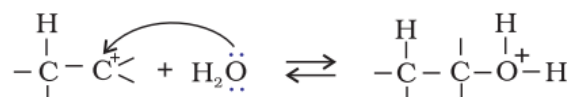
28.	<p>a) Lead storage battery is a secondary battery.</p> <p>Anode: $\text{Pb(s)} + \text{SO}_4^{2-}(\text{aq}) \rightarrow \text{PbSO}_4(\text{s}) + 2\text{e}^-$</p> <p>Cathode: $\text{PbO}_2(\text{s}) + \text{SO}_4^{2-}(\text{aq}) + 4\text{H}^+(\text{aq}) + 2\text{e}^- \rightarrow \text{PbSO}_4(\text{s}) + 2\text{H}_2\text{O(l)}$</p> <p>Overall reaction: $\text{Pb(s)} + \text{PbO}_2(\text{s}) + 2\text{H}_2\text{SO}_4(\text{aq}) \rightarrow 2\text{PbSO}_4(\text{s}) + 2\text{H}_2\text{O(l)}$</p> <p>b) Ions are not involved in the overall cell reaction in the mercury cell.</p>	<p>1/2</p> <p>1/2</p> <p>1/2</p> <p>1/2</p> <p>1</p>
29.	<p>a) i) CH_3Br</p> <p>ii) $\text{CH}_3\text{CH}_2\text{Cl}$</p> <p>b)</p> <p>$\text{CH}_3\text{-CH}_2\text{-CH=CH}_2 + \text{HBr} \xrightarrow{\text{Peroxide}}$ (Anti - Markovnikovs reaction)</p> <p>But - 1 - ene</p> <p>$\text{CH}_3\text{-CH}_2\text{-CH}_2\text{-CH}_2\text{-I} \xleftarrow[\text{dry}]{\text{NaI}}$ $\text{CH}_3\text{-CH}_2\text{-CH}_2\text{-CH}_2\text{-Br}$</p> <p>1 - Iodobutane acetone 1 - Bromobutane</p>	<p>1</p> <p>1</p> <p>1</p>
30.	<p>a) Carbon-oxygen bond in phenol has a partial double bond character due to resonance.</p> <p>b) It is due to symmetry of para-isomers that fits in crystal lattice better as compared to ortho- and meta-isomers.</p> <p>c) Grignard reagents are highly reactive and react with any source of proton to give hydrocarbons.</p> <p>d) Due to -I effect of halogen, ring is deactivated. (Any 3 correct answers)</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p>
31.	<p>a) E and F</p> <p>b) A and B</p> <p>c) Exothermic reaction.</p> <p>The intermolecular attractive forces between C and D is stronger than those Between C-C and D-D. Therefore, energy is released.</p> <p style="text-align: center;">OR</p> <p>c) $P_{\text{total}} = P_1^0 + (P_2^0 - P_1^0) X_2$</p> <p>$400 = 350 + (500 - 350) X_2$</p> <p>$X_2 = 1/3 = 0.33$</p> <p>$X_1 = 2/3 = 0.67$</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1/2</p> <p>1/2</p> <p>1/2</p> <p>1/2</p>
32.	<p>a) Phosphodiester linkage.</p> <p>b) The two strands in DNA are complementary to each other because the hydrogen bonds are formed between specific pairs of bases.</p> <p>c) DNA - Adenine (A), guanine (G), cytosine (C) and thymine (T).</p> <p>RNA - Adenine (A), guanine (G), cytosine (C) and uracil (U).</p> <p style="text-align: center;">OR</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p>

34.

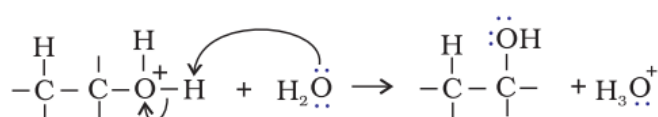
- a) Step 1: Protonation of alkene to form carbocation by electrophilic attack of H_3O^+ .



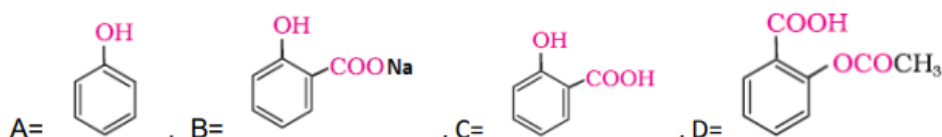
- Step 2: Nucleophilic attack of water on carbocation.



- Step 3: Deprotonation to form an alcohol.

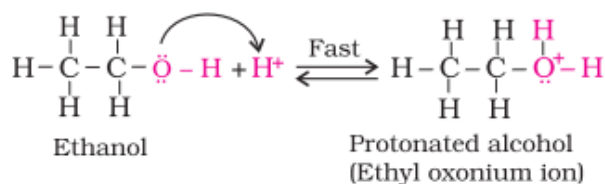


- b)

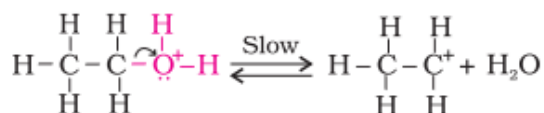


OR

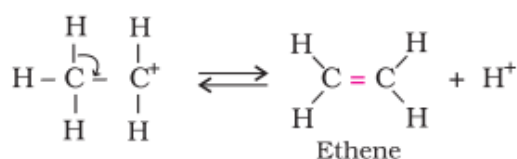
- a) Step 1: Formation of protonated alcohol.



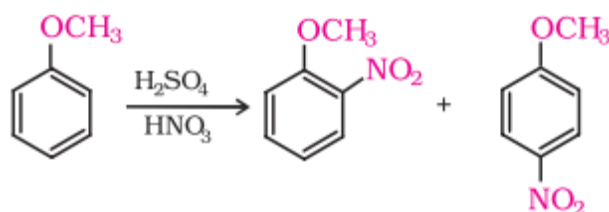
- Step 2: Formation of carbocation: It is the slowest step and hence, the rate determining step of the reaction.



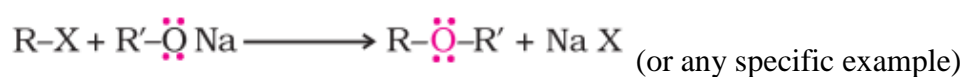
- Step 3: Formation of ethene by elimination of a proton.



- b) i)



- ii)



<p>35.</p>	<p>a) i) Oxygen and fluorine have small size and high electronegativity. Hence, they can oxidize the metal to highest oxidation states.</p> <p>ii) This is because small atoms like B, C, H, N etc. can occupy interstitial sites in the lattice of transition elements.</p> <p>iii) This is because 5f electrons in actinoids have poorer shielding effect than 4f electrons in lanthanoids.</p> <p>b) $5\text{NO}_2^- + 2\text{MnO}_4^- + 6\text{H}^+ \rightarrow 2\text{Mn}^{2+} + 5\text{NO}_3^- + 3\text{H}_2\text{O}$</p> <p>c) $2\text{KMnO}_4 \rightarrow \text{K}_2\text{MnO}_4 + \text{MnO}_2 + \text{O}_2$</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>
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