16-08-15_Sr. IPLCO_JEE-ADV_(2012_P1)_RPTA-3_Q'Paper

JEE-ADVANCED-2012-P1-Model

Time: 3:00 Hrs.

IMPORTANT INSTRUCTIONS

Max Marks: 210

PHYSICS:

Section	Question Type	+Ve Marks	- Ve Marks	No.of Qs	Total marks
Sec – I(Q.N : 1 – 10)	Questions with Single Correct Choice	3	-1	10	30
Sec – II(Q.N : 11 – 15)	Questions with Multiple Correct Choice	4	0	5	20
Sec – III(Q.N : 16 – 20)	Questions with Integer Answer Type	4	0	5	20
Total			20	70	

CHEMISTRY:

Section	Question Type	+Ve Marks	- Ve Marks	No.of Qs	Total marks
Sec – I(Q.N : 21 – 30)	Questions with Single Correct Choice	3	-1	10	30
Sec – II(Q.N : 31 – 35)	Questions with Multiple Correct Choice	4	0	5	20
Sec – III(Q.N : 36 – 40)	Questions with Integer Answer Type	4	0	5	20
Total			20	70	

MATHEMATICS:

Section	Question Type	+Ve Marks	- Ve Marks	No.of Qs	Total marks
Sec - I(Q.N : 41 - 50)	Questions with Single Correct Choice	3	-1 -	10	30
Sec – II(Q.N : 51 – 55)	Questions with Multiple Correct Choice	4	0	5	20
Sec – III(Q.N : 56 – 60)	Questions with Integer Answer Type	4	0	5	20
Total			20	70	

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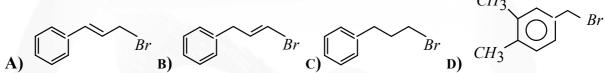
CHEMISTRY

Max.Marks:70

SECTION – I (SINGLE CORRECT CHOICE TYPE)

This section contains 10 multiple choice questions. Each question has 4 choices (A), (B), (C) and (D) for its answer, out of which ONLY ONE is correct

21. Which of the following does NOT undergo SN^2 displacement readily?



22. What nucleophile could be used to react with 1, 4 – dibromobutane to prepare the following compound (You may assume that base is present to remove excess protons)

$$N-CH_3$$
A) CH_3-NH_2 **B)** NH_3 **C)** $(CH_3)_2 NH$ **D)** $N-H$

- **23.** Reaction of (R)-2-chloro-4-methylpentane with excess NaI in acetone gives racemic 2-iodo-4-methylpentane. How can this be explained?
 - A) the reaction mechanism changes to S_N1
 - B) the reaction proceeds via a rapidly inverting radical intermediate
 - C) the substitution is S_N2 , but repeated attack by iodide anion (with inversion) leads to racemisation
 - **D)** iodide anion preferentially attacks chlorine, giving a rapidly inverting carbanion intermediate

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24. Consider the S_N1 solvolysis of the following 1°-alkyl chlorides in aqueous ethanol.

I. $CH_3CH_2CH_2Cl$ II $CH_2 = CHCH_2Cl$ III. CH_3OCH_2Cl IV. $CF_3CF_2CH_2Cl$

What is the order of decreasing reactivity?

A) III > II > IV

B) |I| > I > |I| > IV

C) IV > III > II > I

- $\mathbf{D}) \mathbf{I} > \mathbf{II} > \mathbf{III} > \mathbf{IV}$
- 25. Stereoisomers I and II undergo E2 elimination on treatment with sodium ethoxide in ethanol One isomer reacts 500 times faster than the other. Also, one isomer gives X as the only product, whereas the other gives Y together with some X. Which of the following statements provides the best assignment of I and II?

- A) II reacts faster and gives both Y & X
- B) II reacts faster and gives only X
- c) I reacts faster and gives both Y & X
- D) I reacts faster and gives only X
- **26.** If the rate of reaction of [0.1 M] sodium cyanide with [0.1M] 1-bromoethane is

 1.4×10^{-4} . What effect will an increase in *NaCN* concentration to [0.3] and alkyl bromide concentration to [0.2] have on the overall reaction rate?

- A) increase by 2 times
- B) increase by 3 times
- c) increase by 6 times
- D) increase by 1.5 times

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- Which of the following organic chlorides will not give a Friedel-Craft alkylation product when heated with benzene and AlCl3 (Consider Alkenylation and Alkynylation as alkylation if any in the given reaction)
 - A) $(CH_3)_3CCl$
- B) $CH_2 = CHCH_2Cl$ C) CH_3CH_2Cl D) $CH_2 = CHCl$
- Which reactants combine to give the species shown below as a reactive intermediate? 28.

$$\begin{array}{c} CH(CH_3)_2 \\ H \\ H \\ H \\ \end{array}$$

- A) Benzene, isopropyl bromide, and HBr
- B) Bromobenzene, isopropyl chloride, and AlCl₃
- c) Isopropylbenzene, Br2 and FeBr3
- D) Isopropylbenzene, Br_2 , light, and heat

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29. Which of the following reaction sequences would be best for converting para-

bromoanisole to ortho-ethylanisole?

$$\begin{array}{ccc}
OCH_3 & OCH_3 \\
& ? & & & & & & & \\
Br & & & ? & & & & & & & \\
\end{array}$$

- A) (i) $H_2 \& Pt$ catalyst (ii) $C_2H_5Cl \& AlCl_3$
- B) (i) Mg in ether (ii) aqueous alcohol (iii) $C_2H_5Cl \& AlCl_3$
- C) (i) Mg in ether (ii) $C_2H_5Cl \& AlCl_3$
- **D)** (i) $C_2H_5Cl \& AlCl_3$ (ii) Mg in ether (iii) aqueous alcohol
- **30.** Which reagents are needed to complete the transformation given below?

$$\bigcirc CH_3 \longrightarrow \bigcirc$$

A) $SOCl_2$, $AlCl_3$ B) NBS, Aq NaOH C) Cl_2 / hv , $AlCl_3$ D) $Cl_2 / AlCl_3$

SECTION – II (MULTIPLE CORRECT CHOICE TYPE)

This section contains 5 multiple choice questions. Each question has 4 choices (A), (B), (C) and (D) for its answer, out of which ONE OR MORE is / are correct

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31. Select better combination of reactants which gives following compound as major product in EAS reaction in presence of suitable catalyst

$$H_3C$$
 H_3C
 H_3C

32. Orientation in EAS reactions is predicted based on relative stability of intermediate arenium ion..Apply this concept and identify which of following reactions are correct w.r.t to orientation?

A)
$$\stackrel{\text{H}_2SO_4}{\stackrel{\text{N}}{\longrightarrow}}$$
 $\stackrel{\text{Br}}{\stackrel{\text{N}}{\longrightarrow}}$ $\stackrel{\text{Br}}{\stackrel{\text{N}}{\longrightarrow}}$ $\stackrel{\text{Br}}{\stackrel{\text{N}}{\longrightarrow}}$ $\stackrel{\text{Br}}{\stackrel{\text{N}}{\longrightarrow}}$ $\stackrel{\text{H}_2SO_4}{\stackrel{\text{N}}{\longrightarrow}}$ $\stackrel{\text{N}}{\stackrel{\text{N}}{\longrightarrow}}$ \stackrel

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33. Below are two potential methods for preparing the same ether but only one of them is successful. Identify the successful approach and choose correct choice(s) to explain your method.

$$(I) \xrightarrow{I} CH_3ONa \xrightarrow{OCH_3}$$

$$CH_3 - I \xrightarrow{(CH_3)_3 CONa} OCH_3$$

- **A)** (I): $S_N 1$ is favoured in (I)
- **B)** (II) : $E_1 \& E_2$ are not favoured in (II)
- C) (I): E_2 is not favoured in (I)
- **D)** (II) : $S_N 2$ is most favoured
- 34. Choose the correct order(s):

$$\begin{array}{c}
O \\
\parallel \\
A) & CH_3 - C - CH_2Br > CH_2 = CH - CH_2Br & (S_N1)
\end{array}$$

- B) $CH_3OCH_2Cl > CH_3CH_2Cl (S_N1)$
- C) $CF_3SO_3^- > CH_3SO_3^-$ (Leaving ability)
- **D)** $Et S CH_2CH_2Cl > Et O CH_2 CH_2 Cl$ (Rate of hydrolysis)

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35. Which of the following are correct order of nucleophilicity?

$$(CH_3CH_2)_3N >$$

A)

B)
$$H - O - O^- > H - O^-$$

C) $I^- > F^-(polar\ protic\ solvent)$

$$CH_3O$$
 CH_3O CH_3-C CH_3-C COO^-

SECTION -III (INTEGER ANSWER TYPE)

This section contains 5 questions. The answer to each of the questions is a single digit integer, ranging from 0 to 9. The appropriate bubbles below the respective question numbers in the ORS have to be darkened.

36.

$$\begin{array}{c}
D_2SO_4 \\
D_2O
\end{array}$$

Molecular mass of product is higher than that of benzene by _____units

(Consider excess reagent)

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37. Each of following compounds contain two aromatic (benzene) rings. In every case two rings are different and labeled as A and B. Identify the number of compounds in which EAS reaction is more favoured in ring A compared to ring B

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38. Identify the number of compounds which can't be made using a direct Friedel-Crafts alkylation as major product. (it is necessary to perform an acylation followed by a Clemmensen reduction to avoid carbocation rearrangements and some may not obtained even by this process)

39. Following isomerisation occurs by how many steps?

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40. There are certain factors which favours 'SN' reactions to follow either S_N1 or S_N2 path, structure of substrate is one of them. In how many of following pairs of substrates, structure of first member is more favoured for S_N2 compared to next one?

$$(a) \xrightarrow{CI} \qquad (b) \xrightarrow{Br} \xrightarrow{Br} \qquad (c)$$

$$(c) \xrightarrow{CI} \qquad (d) \xrightarrow{CI} \qquad (d) \xrightarrow{CI} \qquad (e) \xrightarrow{CI} \qquad (f) \xrightarrow{Br} \qquad Br$$

$$(g) \xrightarrow{CI} \qquad (h) \xrightarrow{Br} \qquad I$$

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