Sri Chaitanya IIT Academy

16-08-15_Sr.IPLCO_JEE-ADV_(2012_P1)_RPTA-3_Key&Sol's



Sri Chaitanya IIT Academy, India

A.P, TELANGANA, KARNATAKA, TAMILNADU, MAHARASHTRA, DELHI, RANCHI A right Choice for the Real Aspirant

ICON CENTRAL OFFICE, MADHAPUR-HYD

 Sec: Sr.IPLCO
 JEE-ADVANCE
 Date: 16-08-15

 Time: 3 Hours
 2012-P1-Model
 Max Marks: 210

PAPER-I KEY & SOLUTIONS

PHYSICS

1	D	2	В	3	A	4	В	5	D	6	С
7	С	8	A	9	В	10	D	11	ABCD	12	AB
13	BD	14	ACD	15	AB	16	1	17	1	18	6
19	7	20	4								

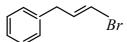
CHEMISTRY

21	В	22	A	23	С	24	A	25	A	26	С
27	D	28	C	29	D	30	С	31	BD	32	ABCD
33	BD	34	BCD	35	BCD	36	6	37	5	38	4
39	4	40	2								

MATHS

41	D	42	В	43	D	44	В	45	C	46	В
47	В	48	C	49	C	50	D	51	A,B,C,D	52	A,C
53	В,С	54	A,B,C	55	A,B,C,D	56	8	57	0	58	1
59	4	60	2								

CHEMISTRY



- 21. Vinyl halides are less reactive in 'SN2' reactions.
- is vinylic halide.

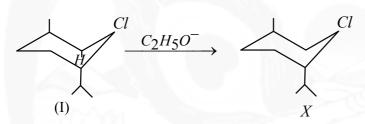
22.
$$Br - CH_2 - CH_2 - CH_2 - CH_2 - Br \xrightarrow{CH_3 - NH_2} \xrightarrow{-H^+}$$

$$Br - CH_2 - CH_2 - CH_2 - CH_2 - NH_2 \xrightarrow{(SN)internal} N - CH_3$$

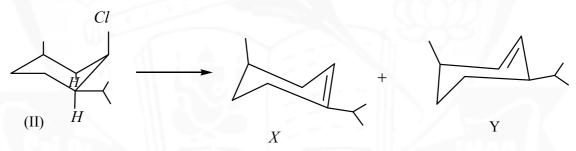
23. R-2-Chloro-4-methyl pentane $\xrightarrow{NaI/acetone}$ S-2-Iodo-4-methyl pentane

$$\underbrace{\frac{NaI/acetone \, S_N \, 2}{R-2-Iodo-4-Methyl}}$$
 R - 2 - Iodo - 4 - Methyl pentane

- 24. Reactivity in sn1 solvolosys increses with increse in stability of intermediate carbocation.
- 25.



I is O less stable conformer, so rate of reaction is less



- 26. Give reaction follows S_N 2 meachanism
 - $\therefore r = K[BrCH_2 CH_3][NaCN]$

As [NaCN] is doubled and $[BrCH_2 - CH_3]$ is tripled so rate is increased by 6 times

- 27. Vinyl chloride are not suitable halogen derivatives for Friedal-Crafts alkylation
- 28. It is one of resonance structure of arenium ion formed by attack of Br^{+2} at para position of isopropyl benzene.
- 29.

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$$CH_3 \xrightarrow{Cl_2} hv \longrightarrow CH_2Cl \xrightarrow{AlCl_3} \longrightarrow$$

30.

31. The combination that follows is not effective, because it involves a Friedel–Crafts reaction on a deactivated aromatic ring

$$H_3C$$
 — CCl + $AICl_3$ no reaction HO_2C

p-Methylbenzoyl chloride

Benzoic acid

The following combination, utilizing toluene, therefore seems appropriate:

$$H_3C$$
 \longrightarrow H_3C \longrightarrow H_3C \longrightarrow HO_2C

32. Sulfonation of furan takes place at C-2. The cationic intermediate is more stable than the cyclo- hexadienyl cation formed from benzene because it is stabilized by electron release from oxygen.

Pyridine reacts with electrophiles at C-3. It is less reactive than benzene, and the carbocation intermediate is less stable than the corresponding intermediate formed from benzene

C-1 of naphthalene is more reactive than C-2 toward electrophilic aromatic substitution.

Naphthalene

33.

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$$CH_{3}ONa \longrightarrow (Major)$$

$$CH_{3}-I \xrightarrow{(CH_{3})_{3}CONa} \xrightarrow{OCH_{3}} (Major)$$

- 34. A, B S_N 1 is more favorable, if carbocation is more stable.
 - C Weak base is good leaving group
 - D Sulpher atom act as good neighboring group
- 35. Conceptual

36.
$$C_6H_6 \xrightarrow{D_2SO_4} C_6D_6$$

37. In the following compound ring A is more reactive

$$A \longrightarrow B \longrightarrow A \longrightarrow NH \longrightarrow B$$

$$A \longrightarrow NH \longrightarrow$$

38. Carbocation is electrophile in Fridel-Crafts alkylation, imagine structure of carbocation has to attack to get given product. If it is the cation which does not rearrange, given compound is major product of Fridel-Crafts alkylation.

$$\begin{array}{c|c}
OH & & & & \\
\hline
OH_2 & & & \\
\hline
& & &$$

$$OH \qquad OH_2 \qquad H_2O \qquad H_$$

39.