DPP - Daily Practice Problems

Chapter-wise Sheets

Date :	Start Time :	End Time :	,
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CHEMISTRY (CC25)

SYLLABUS: Alcohols, Phenols and Ethers

Max. Marks: 180 Marking Scheme: + 4 for correct & (-1) for incorrect Time: 60 min.

INSTRUCTIONS: This Daily Practice Problem Sheet contains 45 MCQ's. For each question only one option is correct. Darken the correct circle/ bubble in the Response Grid provided on each page.

- 1. Diethyl ether reacts, inspite of its usual inert nature, with:
 - (a) Dilute suphuric acid
 - (b) Dilute sodium hydroxide
 - (c) Boron trifluoride
 - (d) Metallic sodium
- 2. *n*-Propyl alcohol and isopropyl alcohol can be chemically distinguished by which reagent?
 - (a) PCI₅
 - (b) Reduction
 - (c) Oxidation with potassium dichromate
 - (d) Ozonolysis
- 3. Which of the following reactions will not result in the formation of anisole?
 - (a) Phenol + dimethyl sulphate in presence of a hase
 - (b) Sodium phenoxide is treated with methyl iodide

- (c) Reaction of diazomethane with phenol
- (d) Reaction of methylmagnesiun iodide with phenol
- 4. Intermolecular hydrogen bonding is strongest in:
 - (a) Methylamine
- (b) Phenol
- (c) Formaldchyde
- (d) Methanol
- 5. Vinyl carbinol is
 - (a) $HO-CH_2-CH=CH_2$
 - (b) $CH_3C(OH) = CH_2$
 - (c) $CH_3 CH = CH OH$
 - (d) $CH_3 C(CH_2OH) = CH_2$
- 6. Lucas reagent is
 - (a) Cone, HCl and anhydrous ZnCl₂
 - (b) Cone. HNO₃ and hydrous ZnCl₂
 - (c) Conc. HCl and hydrous ZnCl₂
 - (d) Conc. HNO₃ and anhydrous ZnCl₂

RESPONSE GRID 1. abcd 6. abcd

2. (a)(b)(c)(d)

3. (a)(b)(c)(d)

4. (a)(b)(c)(d)

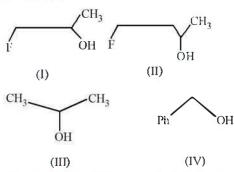
5. (a)(b)(c)(d)

Space for Rough Work .

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DPP/CC25

The order of reactivity of the following alcohols towards conc. HCl is

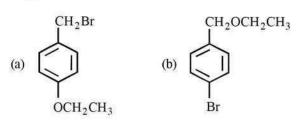


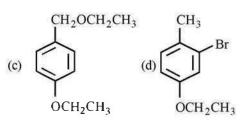
- (a) I>II>III>IV(b)
- I>|||>|V
- (c) IV > III > II > I(d)
- [V>[[>]>]
- What is the major product M in the following reaction?

$$CH_3$$

$$NBS,benz \bullet ylpcroxide \longrightarrow X \xrightarrow{CH_3CH_2ON_3} M$$

$$CCl_4,heat \longrightarrow CII_3CH_2OII$$





- 9. Epichlorohydrin is
 - (a) 3-Chloropropane
 - (b) 3-Chloropropan-I-ol
 - (c) 3-Chloro-1, 2-cpoxypropane
 - (d) None of these
- 10. CH₂CH₂OH can be converted into CH₂CHO by
 - (a) catalytic hydrogenation
 - (b) treatment with LiAlH,

- (c) treatment with pyridinium chlorochromate
- (d) treatment with KMnO₄
 11. In Williamson synthesis if tertiary alkyl halide is used than
 - (a) other is obtained in good yield
 - (b) ether is obtained in poor yield
 - (c) alkene is the only reaction product
 - (d) a mixture of alkene as a major product and ether as a minor product forms.
- 12. Denaturation of alcohol is the
 - mixing of CuSO₄ (a foul smelling solid) and pyridine (to give the colour) to make the commercial alcohol unfit for drinking
 - mixing of CuSO₄ (to give the colour) and pyridine (a foul smelling solid) to make the commercial alcohol unfit for drinking
 - mixing of Cu(OAc), and ammonia to make the commercial alcohol unfit for drinking
 - mixing of Cu(OAc), and pyridine to make the commercial alcohol unfit for drinking
- 13. 2-Phenylethanol may be prepared by the reaction of phenylmagnesium bromide with
 - (a) HCHO
- (b) СН,СНО
- (c) CH₂COCH₂
- (d)
- 14. Arrange the following in increasing order of their acidity? o-cresol(a), salicyclic acid(b), phenol(c)
 - (a) c < a < b (b) b < c < a
 - (c) a < b < a (d) a < c < b
- Which of the following is most reactive towards aqueous HBr?
 - I-Phenyl-I-propanol
 - (b) I-Phenyl-2-propanol
 - 3-Phenyl-I-propanol
 - All are equally reactive
- The ionization constant of phenol is higher than that of ethanol because:
 - phenoxide ion is bulkier than ethoxide (a)
 - phenoxide ion is stronger base than ethoxide
 - phenoxide ion is stabilized through delocalization
 - (d) phenoxide ion is less stable than ethoxide
- 17. Rectified spirit is a mixture of
 - (a) 95% ethyl alcohol + 5% water
 - 94% ethyl alcohol +4.53 water (b)
 - 94.4% ethyl alcohol + 5.43% water (c)
 - 95.87% ethyl alcohol +4.13% water

RESPONSE GRID

- 7. (a)(b)(c)(d)
- **8.** (a)(b)(c)(d)
- 9. (a)(b)(c)(d)
- 10.(a)(b)(c)(d)

- 12.abcd 17.abcd
- 13.(a)(b)©(d)
- 14. (a) (b) (c) (d)
- 15.(a)(b)(c)(d)

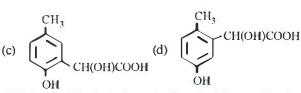
- 18. Ethanol is prepared industrially by
 - hydration of ethylene (b) fermentation of sugar
 - Both the above
- (d) None of these
- 19. Mechanism of acid catalysed hydration reaction involves Protonation of alkene to form carbocation by electrophilic attack of H2O+
 - Nucleophilic attack of water on carbocation.
 - (iii) Deprotonation to form alcohol.
 - (a) (i)and(ii)
- (b) (i)and(iii)
- (c) (i), (ii) and(iii)
- (ii) and (iii) (d)
- 20. Match the columns

Column-I

Column-II

- Methanol
- Conversion of phenol to o-hydroxysalicylic acid
- B. Kolhe's reaction Π. Wood spirit
- C. Heated copper at 573 K Williamson's synthesis III.
- Conversion of 2° Reaction of alkyl halide D IV alcohol to ketone with sodium alkoxide
- (a) A-IV; B-III; C-II; D-1
- (b) A-II; B-IV; C-I; D-III
- (c) A-II; B-I; C-IV; D-III
- (d) A III; B II; C I; D IV
- 21. Absolute alcohol (100% alcohol) is prepared by distilling rectified spirit over
 - (a) Na
- (b) CaCl₃
- (c) Mg
- (d) $Mg(\bar{O}C_2H_5)$,
- 22. p-cresol reacts with chloroform in alkaline medium to give the compound A which adds hydrogen cyanide to form, the compound B. The latter on acidic hydrolysis gives chiral carboxylic acid. The structure of the carboxylic acid is

(a)
$$CH_3$$
 (b) CH_2COOH OH



- Which one of the following will show the highest pH value?
 - (a) m-nitrophenol.
- (b) p-nitrophenol.
- (c) o-nitrophenol.
- (d) Both (b) and (c).

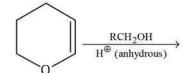
- Which of the following compounds is resistant to nucleophilic attack by hydroxyl ions?
 - (a) Methyl acetate
- (b) Acetonitrile
- Acetamide
- (d) Diethyl ether
- Zerevitinov's determination of active hydrogen in a compound is based upon its reaction with
 - (a) Na
- (b) CH₂Mgl
- (c) Zn
- (d) Al
- Williamson's synthesis is used to prepare
 - (a) acctone
- (b) diethylether
- (c) P.V.C.
- (d) bakelite
- Which of the following statements are correct?
 - (i) Ethanol mixed with methanol is called denatured alcohol.
 - (ii) Excess of methanol in body may cause blindness.
 - (iii) In the body methanol is oxidised to methanoic acid.
 - (iv) A methanol poisoned patient is treated by giving intravenous injections of ethanoic acid.
 - (a) (i),(ii) and(iii)
- (b) (ii), (iii) and(iv)
- (c) (i) and (v)
- (d) (i), (iii) and (iv)
- 28. In the following sequence of reactions,

$$CH_3CH_2OH \xrightarrow{P+I_2} \Lambda \xrightarrow{Mg} B \xrightarrow{HCHO}$$

$$C \xrightarrow{H_2O} D$$

the compound D is

- (a) propanal
- (b) butanal
- (c) *n*-butyl alcohol
- (d) n-propyl alcohol
- When wine is put in air, it becomes sour due to
 - (a) hacteria
 - (b) oxidation of C₂H₅OH to CH₃COOH
 - (c) virus
 - (d) formic acid formation
- Which of the following diols would cleave into two fragments with HIO₄
 - (a) 1, 3-hexanediol
- (b) 2,4-hexanediol
- (c) 1,6-hexanediol
- (d) 3, 4-hexanediol
- 31. The major product of the following reaction is



- (a) a hemiacetal
- **(**b) an acetal
- (c) an ether
- (d) an ester

RESPONSE GRID

- 18. (a) (b) (c) (d) 23.(a)(b)(c)(d)
- 19.(a)(b)(c)(d) 24.(a)(b)(c)(d)
- **20.**(a)(b)(c)(d) 25.(a)(b)(c)(d)
- 21. (a) (b) (c) (d) 26.(a)(b)(c)(d)
- **22.** (a)(b)(c)(d) 27. (a)(b)(c)(d)

- 28.(a)(b)(c)(d) 29.(a)(b)(c)(d)
- **30.**(a)(b)(c)(d)
- 31.(a)(b)(c)(d)

C-100 DPP/ CC25

32. H₂COH · CH₂OH on heating with periodic acid gives:

(a) 2 HCOOH

(c) 2 H = O

- (d) 2 CO₂
- 33. Victor Meyer's test is not given by
 - (a) (CH₃)₃COH
- (b) C₂H₅OH
- (c) (CH₁), CHOH
- (d) CH,CH,CH,OH
- 34. What is X in the following reaction?

$$CH_{2} \xrightarrow{O} C-CH_{3} \xrightarrow{X} H_{2}C-C-CH_{3}$$

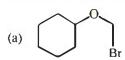
$$CH_{3} \xrightarrow{CH_{3}} CH_{3}$$

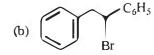
- (a) CH₃OH,H₂SO₄
- (b) CH₃OH, CH₃O⁻ Na
- (c) H₂O/H₂SO₄ followed by CH₃OH
- (d) CH₃MgBr/cther followed by H₃O⁺
- 35. Which of the following pairs of reagents would give 4-methyl-2-pentanol?
 - (a) (CH₃)₂CHLi,CH₃COCH₃
 - (b) (CH₃)₂CHCH₂Li,CH₃CHO
 - (c) (CH₃)₂CHLi,CH₃CH₂CHO
 - (d) CH₃CH₂Li₂(CH₃)₂CH.CHO
- **36.** Which of the following cannot be made by reduction of ketone or aldehyde with NaBH₄ in methanol?
 - (a) I butanol
- (b) 2 butanol
- (c) 2 methyl 1 propanol (d) 2 methyl 2 propanol
- 37. Osmium tetraoxide is a reagent used for
 - (a) hydroxylation of acetylenes
 - (b) hydroxylation of olefins to give cis-diols
 - (c) hydroxylation of olefins to form trans-diols
 - (d) hydroxylation of carbonyl compounds
- **38.** The reaction of sodium ethoxide with ethyl iodide to form diethyl other is termed
 - (a) electrophilic substitution
 - (b) nucleophilic substitution
 - (c) electrophilic addition
 - (d) radical substitution

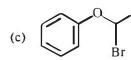
- 39. The IUPAC name of $CH_3 CH CH_2 C CH_3$ is OH OH
 - (a) 1, 1-dimethyl-1, 3-butanediol
 - (b) 2-methyl-2, 4-pentanediol
 - (c) 4-methyl-2, 4-pentanediol
 - (d) 1, 3, 3-trimethyl-1, 3-propanediol
- 40. Give IUPAC name of the compound given below

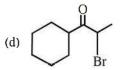
$$\begin{array}{cccc} \operatorname{CH_3} - \operatorname{CH} - \operatorname{CH_2} - \operatorname{CH_2} - \operatorname{CH} - \operatorname{CH_3} \\ & & \operatorname{OH} \end{array}$$

- (a) 2-Chloro-5-hydroxyhexane
- (b) 2-Hydroxy-5-chlorohexane
- (c) 5-Chlorohexanc-2-ol
- (d) 2-Chlorohexan-5-ol
- 41. Aspirin is an acctylation product of
 - (a) p-Dihydroxybenzene (b) o-Hydroxybenzoic acid
 - (c) o-Dihydroxybenzene (d) m-Hydroxybenzoic acid
- **42.** Acetic anhydride reacts with diethyl ether in the presence of anhydrous AlCl₃ to give
 - (a) CH, COOCH,
- (b) CH₃CH₂COOCH₃
- (c) CH, COOCH, CH,
- (d) CH₃CH₂OH
- 43. Formation of which compound given below from 1-butanol needs an oxidising agent?
 - (a) $CH_3CH_2CH_2CH_2Br$ (b) $CH_3CH_2CH_2CH = O$
 - c) $(CH_3CH_2CH_2CH_2)_2 \bullet (d) CH_3 CH_2CH = CH_2$
- 44. $o-Xylene \xrightarrow{HNO_3} X \xrightarrow{Phenol} Y$. The product Y is
 - (a) Phthalicacid
- (b) Isophthalic acid
- (c) Phenolphthalein
- (d) o-Hydroxysulphonic acid
- 45. Which of the following, upon treatment with tert-BuONa followed by addition of bromine water, fails to decolourize the colour of bromine?









RESPONSE GRID

- 32.abcd
- 33.abcd
- **34.**ⓐⓑⓒⓓ
- 35.abcd 40.abcd
 - 36. abcd 41. abcd

- 37.(a)(b)(c)(d) 38.(a)(b)(c)(d) 42.(a)(b)(c)(d) 43.(a)(b)(c)(d)
- 39.abcd 44.abcd
 - 45.abcd
-) 41.

DAILY PRACTICE PROBLEMS

CHEMISTRY SOLUTIONS

DPP/CC25

- (c) Boron trifluoride being a lewis acid forms adduct with diethyl ether which is a lewis base.
- 2. (c) Primary alcohol on oxidation give aldehyde which on further oxidation give carboxylic acid whereas secondary alcohols give ketone.

$$CH_3CH_2CHO \xrightarrow{[O]} CH_3CH_2COOH$$

$$\frac{H_3C}{H_3C} > CH - OH \xrightarrow{[O]} \frac{H_3C}{H_3C} > C = O$$
Ketono

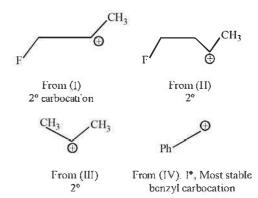
 (d) Phenol has active (acidic) hydrogen so it reacts with CH₃MgI to give CH₄, and not anisole

$$C_6H_5OH + CH_3MgI \longrightarrow CH_4 + C_6H_5OMgI$$

4. (d) Hydrogen bonding is formed in compounds in which H is attached to highly electronegative element like F, O and N.

O []
(i) In
$$H - C - H$$
, \bullet is not having H atom so it shows very little H-bonding.

- (ii) N is less electronegative than ●, so H—bond formed by amines will be weak than that by alcohols.
- (iii) C₆H₅OH forms weak H-bonding due to steric hinderance due to bulky phenyl group.
- 5. (a) Methyl alcohol (CH_3OH) is also known as carbinol. Hence vinyl carbinol is $CH_2 = CH - CH_2OH$
- 6. (a) Lucas reagent is cone. HCl + anhyd. ZnCl₂.
- 7. (c) The order of reactivity depends upon the stability of the carbocations formed.



Remember that presence of electron-withdrawing group intensifies i.e., destabilises the carbocation thus (i) and (ii) are less stable than (iii). Further (i) is less stable than (ii) because —I effect is more pronounced in (i) due to less distance between F and + charge. Thus the stability order of the four carbocations and reactivity of their parent alcohols will be

$$\begin{array}{c}
\text{CH}_2\text{OCH}_2\text{CH}_3\\
\hline
\text{CH}_3\text{CH}_2\text{OH}
\end{array}$$

9. (c)
$$CH_2$$
— $CHCH_2CI$ – 3-chloro -1, 2-epoxypropane

- 10. (c)
- 11. (c) If a tertiary alkyl halide is used, an alkene is the only reaction product and no other is formed. For example, the reaction of CH₃ONa with (CH₃)₃C-Br gives exclusively 2-methylpropene.

$$CH_{3}$$

$$CH_{3} - C - Br + \stackrel{+}{N}a\stackrel{-}{\tilde{Q}} - CH_{3} \longrightarrow$$

$$CH_{3} - C = CH_{2} + NaBr + CH_{3} \bullet H$$

$$CH_{3}$$

It is because alkoxides are not only nucleophiles but strong bases as well. They react with alkyl halides leading to elimination reactions.

2-Methylpropenc

12. (b) The commercial alcohol is made unfit for drinking by mixing in it some copper sulphate (to give it colour) and pyridine (a foul smelling liquid). It is known as denaturation of alcohol.

13. (d) 2-Phenylethanol, $CH_2OHCH_2C_6H_5$, is a 1° alcohol which can be prepared from C_6H_5MgBr by treating with ethylene oxide (note that HCHO will introduce only one carbon atom, i.e. it will give $C_6H_5CH_2OH$ and not $C_6H_5CH_2CH_2OH$).

Electron releasing groups (-CH₃, -OCH₃, -NCH₃ etc) intensify the negative charge of phenoxide ion, i.e., destablises it hence decrease ionization of parent phenol. Therefore decreases acidity while electron with drawing groups (-NO₂, -COOH, -CHO etc.) increases acidity.

15. (a) Here also, carbocation is formed as an intermediate, hence the species capable of forming most stable carbocation will be most reactive.

16. (c)
$$C_6H_5OH + H_2O \rightleftharpoons C_6H_5O^- + H_3O^+$$
Phenoxide ion

The phenoxide ion is stable due to resonance.

The negative charge is delocalized in the benzene ring which is a stabilizing factor in the phenoxide ion and because of this reason ionization constant of phenol is

higher whereas no resonance is possible in alkoxide ions (RO⁻) derived from alcohol. The negative charge is localized on oxygen atom in case of alcohols.

- 17. (d)
- 18. (c) Hydration of alkenes

$$CH_2 = CH_2 + HHSO_4 \rightarrow CH_3 - CH_2 - HSO_4$$

 $CH_3 - CH_2HSO_4 \xrightarrow{H_2O} CH_3 - CH_2 - OH + H_2SO_4$

Fernentation of sugar:

$$C_{12}H_{22}O_{11} + H_2O \xrightarrow{\text{Invertase}} C_6H_{12}O_6 + C_6H_{12}O_6$$
Glucose
Fructese

$$C_6H_{12}O_6 \xrightarrow{\text{Nymase}} 2C_2H_5OH + 2CO_2$$
Glucos or Fructose

19. (c) The mechanism of the reaction involves the following three steps:

Step 1: Protonation of alkene to form carbocation by electrophilic attack of H_3O^+

$$H_2O + H^+ \rightarrow H_3O^+$$

$$C = C + H - O + H \rightarrow C + H_2O$$

Step 2: Nucleophilic attack of water on carbocation.

Step 3: Deprotonation to form an alcohol.

$$-\overset{\mid}{\underset{H}{\text{C}}}-\overset{\mid}{\underset{\downarrow}{\text{C}}}-\overset{\mid}{\underset{\downarrow}{\text{C}}}-\overset{\mid}{\underset{\downarrow}{\text{C}}}-\overset{\mid}{\underset{\downarrow}{\text{C}}}+\overset{\mid}{\underset{\downarrow}{\text{C}}}-+\underset{\downarrow}{\text{H}_{3}}\overset{\mid}{\underset{\downarrow}{\text{C}}}$$

20. (c)

21. (d)
$$(C_2H_5O)_2Mg + H_2O \xrightarrow{\text{distill}}$$

 $2C_2H_5OH + Mg(OH)_2$
(distillate) (residue)

22. (c)
$$CH_3$$

$$CHCl_3 + NaOH$$

$$Reimer Tiemman reaction$$

$$CH_3$$

$$CH_3$$

$$CH = \bigoplus_{l} CHCN$$

$$OH H$$

DPP/CC25 — ______ s-71

- 23. (a) In case of m-nitrophenol operational effect of nitrogroup is electron withdrawing inductive effect while in case of b and c, both -R and -l effect are operational.
- **24. (d)** Diethyl ether, being a Lewis base, is not attacked by nucleophiles, while all others contain electrophilic carbon, hence attacked by nucleophiles like OH⁻ ions.

$$\begin{array}{c|c}
O\delta - & \delta + & \delta - \\
CH_3 - C - OCH_3 & CH_3 - C \equiv N
\end{array}$$

$$\begin{array}{c} O_{\delta-} \\ \parallel \\ CH_3-C-NH_2 \end{array} \qquad C_2H_5-O-C_2H_5$$

25. (b) Number of active hydrogen in a compound corresponds to the number of moles of CH₄ evolved per mole of the compound.

$$-NH_2$$
, $-SH$, $-OH$ or $-C \equiv CH \xrightarrow{CH_3Mgl}$
 $-CICH_4 \uparrow (2CH_4 \text{ from } -NH_2)$

- 26. (b) $C_2H_5Br + C_2H_5ONa \xrightarrow{-NaBr} C_2H_5 O C_2H_5$ Sod. ethoxide diethyl ether
- 27. (a) A methanol poisoned patient is treated by giving intravenous injection of ethanol.

28. (d)
$$CH_3CH_2OH \xrightarrow{P+1_2} CH_3CH_2I \xrightarrow{Mg} Ether$$

$$\begin{array}{c} \text{CH}_2\text{CH}_3\\ \text{CH}_3\text{CH}_2\text{MgI} & \xrightarrow{\text{HCHO}} & \text{H-C-OMgI}\\ \text{(B)} & \text{H} \\ \text{(C)} \end{array}$$

29. (b)

30. (d)
$$CH_3 - CH_2 - CH - CH - CH_2 - CH_3 \xrightarrow{HI\Phi_4}$$
OH OH
$$2CH_3 - CH_2 - CHO$$

31. (b)
$$O$$

32. (c) 1, 2 – Diols, when treated with an aqueous solution of periodic acid give aldehyde

$$CH_2OH \xrightarrow{HIO_4} CH_2O+CH_2O$$

Note that a 1° alcohol gives CH₂O. Since in glycol both the OH groups, are primary hence give 2 molecules of CH₂O as by product.

33. (a)

34. (a)
$$CH_2 \xrightarrow{O} C-CH_3 \xrightarrow{CH_3OH} H_2C-C-CH_3 \xrightarrow{CH_3} CH_3$$

35. (b)
$$CH_3 - CH < CH_2 \cdot CH(CH_3)_2 + CH_3 \cdot CHO$$

$$OH \quad CH_3$$

$$CH_3 - CH - CH_2 - CH - CH_3$$

36. (d) 2-methyl-2-propanol is
$$CH_3 - C - CH_3$$
. It cannot be CH_3

obtained by reduction of an aldehyde or ketone with NaBH₄.

37. (b)

38. (b) Reaction of sodium ethoxide with ethyl iodide to produce diethyl ether is known as Williamson synthesis.
It is a nucleophilic substitution reaction and proceeds via S_N2 mechanism.

39. (b)
$$CH_3 - CH - CH_2 - C - CH_3$$

OH OH

2-methyl- 2, 4-pentancdiol.

40. (c)

S-72 DPP/CC25

41. (b)

42. (c) (CH₃CH₂)₂O+(CH₃CO)₂O Ether Acid anhydride

43. (b) $CH_3CH_2CH_2CH_2OH \xrightarrow{Oxidation}$

44. (c)

45. (a)

(a)
$$O$$
Br
 $tert$ -BuONa
 O -tBu

(fails to decolorise the colour of bromine) due to unsaturation

(b)
$$C_6H_5$$
tert-BuONa
(it decolorises bromine solution)

(c)
$$\xrightarrow{\text{Br}} \xrightarrow{\text{tert}-\text{BuONa}} \xrightarrow{\text{O}}$$

(it decolorises bromine solution)

Products formed in option (2), (3) & (4) decolorises bromine solution due to presence of double bond.