



Sample Paper-01

Dropper NEET (2024)

CHEMISTRY

ANSWER KEY

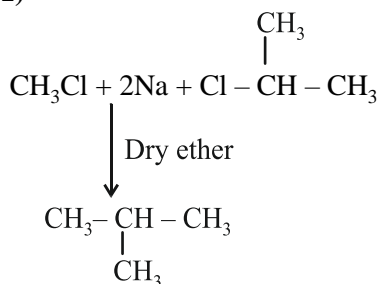
1. (1)
2. (4)
3. (2)
4. (2)
5. (2)
6. (3)
7. (2)
8. (2)
9. (4)
10. (4)
11. (4)
12. (1)
13. (3)
14. (4)
15. (2)
16. (2)
17. (1)
18. (4)
19. (1)
20. (1)
21. (2)
22. (2)
23. (2)
24. (2)
25. (4)

26. (2)
27. (3)
28. (1)
29. (4)
30. (3)
31. (1)
32. (1)
33. (1)
34. (2)
35. (4)
36. (1)
37. (2)
38. (3)
39. (4)
40. (3)
41. (3)
42. (4)
43. (3)
44. (1)
45. (1)
46. (4)
47. (4)
48. (2)
49. (2)
50. (1)

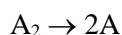


HINTS AND SOLUTION

1. (1)



2. (4)



$$\text{K.E.} = h\nu - h\nu_0$$

$$= 4.4 \times 10^{-19} - 4.0 \times 10^{-19}$$

$$= 0.4 \times 10^{-19} \text{ J}$$

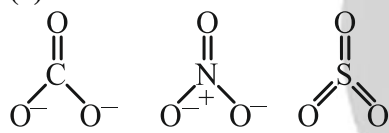
$$= 4 \times 10^{-20} \text{ J}$$

$$\text{K.E. per atom} = 2 \times 10^{-20} \text{ J}$$

3. (2)

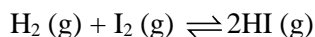
$$\lambda = \frac{12.3}{\sqrt{V}} \text{ \AA}$$

4. (2)



Trigonal planar

5. (2)

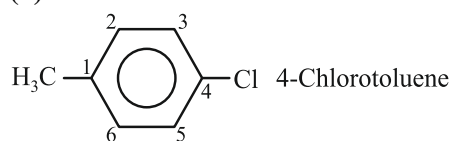


For the given reaction, on increasing temperature K_c decreases, that implies reaction is exothermic.

6. (3)

Addition of electron to O^- is endothermic as O^- will tend to resist the addition of further electron due to inter electronic repulsions.

7. (2)



8. (2)

$$\text{pOH} = \text{pK}_b + \log \frac{[\text{salt}]}{[\text{base}]}$$

$$[\text{NH}_4\text{OH}] = [\text{NH}_4\text{Cl}] = 0.1 \text{ M}$$

$$\text{pOH} = \text{pK}_b$$

$$14 - \text{pH} = \text{pK}_b$$

$$\text{pK}_b = 14 - 9.25 = 4.75$$

9. (4)



(E = C, Si, Ge, Sn, Pb)

Due to decrease in E-H bond strength down the group, reducing nature increases.

10. (4)

Clemmensen reduction

11. (4)

$$n_C = \frac{24}{12} = 2$$

$$n_H = \frac{4}{1} = 4$$

$$n_O = \frac{32}{16} = 2$$

Empirical formula

C:H:O

2:4:2

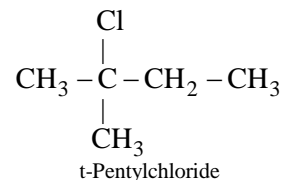
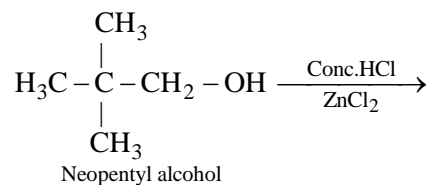
1:2:1

CH_2O

12. (1)

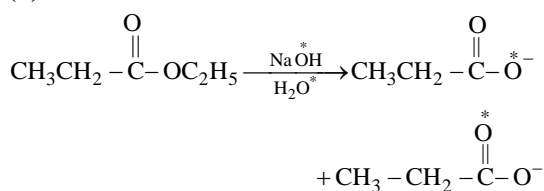
There is no possibility of d-d transition.

13. (3)





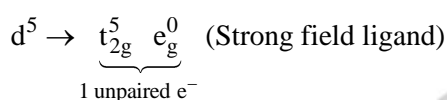
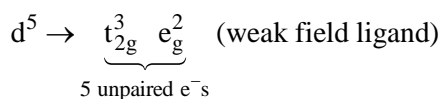
14. (4)



15. (2)

Denaturation disrupts only secondary and tertiary structures of proteins.

16. (2)



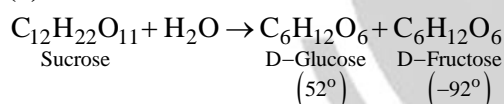
17. (1)

CuSO_4 solution absorbs orange-red colour and appears blue (complimentary to the colour absorbed).

18. (4)

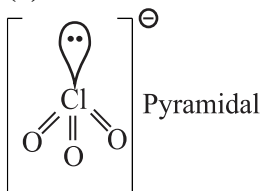
KCl is not present in dry cell.

19. (1)



$$\text{Specific rotation} = \frac{1}{2} \left(52^\circ + (-92^\circ) \right) = -20^\circ$$

20. (1)



21. (2)

$$\frac{K_p}{K_c} = RT^{\Delta n_g}$$

Where Δn_g = number of gaseous mole of product – number of gaseous mole of reactant)

22. (2)

Salt contains Zn^{2+}

23. (2)

$$\begin{aligned} \text{Number of spectral lines} &= \frac{(n_2 - n_1)(n_2 - n_1 + 1)}{2} \\ &= \frac{(7 - 2)(7 - 2 + 1)}{2} = 15 \end{aligned}$$

24. (2)

According to Arrhenius equation, rate of a reaction increases with decrease in activation energy.

25. (4)

Faraday's first law of electrolysis,

$$W = Z \times I \times t$$

where, Z = electrochemical equivalent (g/C)

I = current (Amps)

t = time (sec)

$$W = \frac{63.6}{2 \times 96500} \times 0.5 \times 60 \times 60$$

$$W = 0.56 \text{ g.}$$

26. (2)

$$\text{Solubility of AgCl in water} = \sqrt{K_{sp}} = S_1$$

In 0.01M CaCl_2 , it is given by,

$$K_{sp} = S_2 \times (0.01 \times 2 + S) \left(\because S_2 = \frac{K_{sp}}{0.02} \right)$$

In 0.01 M NaCl , it is given by

$$K_{sp} = S_3 \times (0.01 + S) \left(\because S_3 = \frac{K_{sp}}{0.01} \right)$$

In 0.05M AgNO_3 , it is given by

$$K_{sp} = S_4 \times (0.05 + S) \left(\because S_4 = \frac{K_{sp}}{0.05} \right)$$

The solubilities are derived by neglecting S in comparison to 0.02, 0.01 and 0.05.

So, order of solubilities:

$$S_1 > S_3 > S_2 > S_4$$

27. (3)

The rate constant for first order reaction does not depends on initial concentration of reactants.




28. (1)
A nucleophilic addition reaction is a chemical addition reaction in which a nucleophile forms a sigma bond with an electron-deficient species.

29. (4)
Buffer is a solution of weak acid and conjugate base of weak acid or a weak base and a conjugate acid of weak base, e.g. $\text{NH}_4\text{Cl} + \text{NH}_4\text{OH}$, $\text{CH}_3\text{COOH} + \text{CH}_3\text{COONa}$, $\text{NaCN} + \text{HCN}$

30. (3)
Acetate ion is more resonance stabilized than phenoxide ion.

31. (1)
 $\text{CH} \equiv \text{CH}$, $\text{CH}_2 = \text{CH}_2$, $\text{CH}_3 - \text{CH}_3$
 \uparrow \uparrow \uparrow
 sp sp^2 sp^3
 Electronegativity of carbon atom:
 $\text{sp} > \text{sp}^2 > \text{sp}^3$.
 \therefore Acidic character follows the order
 $\text{CH} \equiv \text{CH} > \text{CH}_2 = \text{CH}_2 > \text{CH}_3 - \text{CH}_3$
 Hence, order of basic strength:
 $\text{CH} \equiv \text{C}^\ominus < \text{CH}_2 = \text{C}^\ominus < \text{CH}_3 - \text{C}^\ominus\text{H}_2$
 (I) (II) (III)

32. (1)
 $\text{CH}_3 - \text{CH}_3 \rightarrow 2 \text{ sp}^3$ carbons
 $\text{CH}_2 = \text{CH}_2 \rightarrow 2 \text{ sp}^2$ carbons
 $\text{CH} \equiv \text{CH} \rightarrow 2 \text{ sp}$ carbons
 $\rightarrow 6 \text{ sp}^2$ carbons

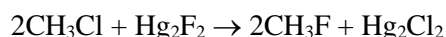
33. (1)
Molecular orbital electronic configuration-
 $(\sigma 1s^2)(\sigma^* 1s^2)(\sigma 2s^2)(\sigma^* 2s^2)(\sigma 2p_z^2)(\pi 2p_x^2)(\pi 2p_y^2)$
 $(\pi^* 2p_x^2)(\pi^* 2p_y^2)$
 Number of electrons in bonding MOs = 10
 Number of electrons in anti-bonding MOs = 8

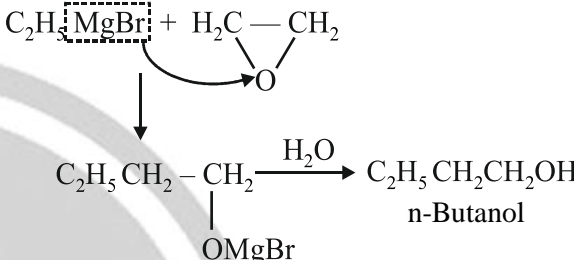
34. (2)
When $T \rightarrow \infty$
then $K = A$

35. (4)
All of the above given statements are correct.

36. (1)
Alkyl fluorides which cannot be prepared by Finkelstein reaction is best accomplished by Swarts reaction. In this reaction, the corresponding fluoride is obtained by the action of mercurous fluoride or antimony trifluoride on corresponding chlorides.

The reaction is as follows:

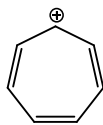


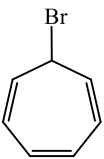
37. (2)


38. (3)
 $\text{Al}_2\text{O}_3 \rightarrow 2\text{Al}^{3+} + 3\text{O}^{2-}$
 $\text{Al}^{3+} + 3\text{e}^- \rightarrow \text{Al}$
 $\therefore 3\text{F charge gives Al} = 27 \text{ g}$
 $\therefore 2\text{F charge will give Al} = \frac{27}{3} \times 2 = 18 \text{ g}$
 [1 mole $\text{e}^- = 1\text{F charge}$]

39. (4)
 $\text{K}_2\text{Cr}_2\text{O}_7 + 7\text{H}_2\text{SO}_4 + 6\text{KI} \rightarrow \text{Cr}_2(\text{SO}_4)_3 + 3\text{I}_2 + 7\text{H}_2\text{O} + 4\text{K}_2\text{SO}_4$

40. (3)
Ammoniacal silver nitrate

41. (3)
Tropylium ion,  is an aromatic species.

Therefore,  will react fastest with AgNO_3 solution.



42. (4)

If A-B interactions > A-A or B-B interactions then mixture shows negative deviation from Raoult's law.

Hexane + Heptane : No deviation

Benzene + Toluene : No deviation

Water + Ethanol : +ve deviation

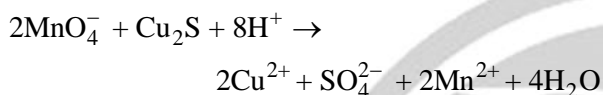
Nitric acid + Water : -ve deviation

43. (3)

Among all monosaccharides and all disaccharides, only sucrose is a non-reducing sugar.

44. (1)

The balanced reaction is:



Thus, the number of moles of MnO_4^- ion that will be needed to oxidise one mole of Cu_2S completely = 2

45. (1)

$$d = M \left(\frac{1}{m} + \frac{\text{molar mass of solute}}{1000} \right)$$

$$d = 11.12 \left(\frac{1}{94.12} + \frac{40}{1000} \right) = 0.556 \text{ g mL}^{-1}$$

46. (4)

Only Terminal alkynes gives white precipitate with ammoniacal silver nitrate solution.

47. (4)

Neutralisation reactions are exothermic in nature.
Acidic strength $\propto \Delta H_{\text{neutralisation}}$ (negative value)

Thus, acidic strength order is:

B > D > A > C

48. (2)



At $t = 0$: 10 m mol 5 m mol 0 0

At equil: 5 m mol 0 5 m mol 5 m mol

Thus, the given buffer is an acidic buffer.

(NaOH is the limiting reagent)

pH of an acidic buffer is given by:

$$\text{pH} = \text{pK}_a + \log \frac{[\text{Salt}]}{[\text{Acid}]}$$

$$= 4.74 + \log \frac{5}{5}$$

$$= 4.74 + \log 1$$

$$= 4.74 + 0$$

$$= 4.74$$

$$[\text{m mol} = \text{Volume} \times \text{Molarity}]$$

49. (2)

$$w = -2.303nRT \log \frac{V_2}{V_1}$$

$$= -2.303 \times 5 \times 8.314 \times 300 \log \frac{60}{6}$$

$$= -28.72 \text{ kJ}$$

50. (1)

The complex exhibits two geometrical isomers (i) facial and (ii) meridional.

