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Answer & Solutions

for

NEET 2022_(Re-Exam)

Chemistry

51. Match List - I with List - II:

(quantum number)

List - I

List- II

` '

(orbital)

(a)
$$n = 2$$
, $\ell = 1$

(i) 2 s

(b)
$$n = 3$$
, $\ell = 2$

(ii) 3 s

(c)
$$n = 3$$
, $\ell = 0$

(iii) 2 p

(d)
$$n = 2$$
, $\ell = 0$

(iv) 3 d

Choose the **correct answer** from the options given below:

Sol. Answer (1)

 $\ell = 0 \Rightarrow \text{s-subshell}$

 $\ell = 1 \Rightarrow p$ -subshell

 ℓ = 2 \Rightarrow d-subshell

 ℓ = 3 \Rightarrow f-subshell

$$\therefore$$
 n = 2, ℓ = 1 \Rightarrow 2 p

$$n = 3$$
, $\ell = 2 \Rightarrow 3 d$

$$n = 3$$
, $\ell = 0 \Rightarrow 3 s$

$$n = 2$$
, $\ell = 0 \Rightarrow 2 s$

- **52.** The density of the solution is 2.15 g mL⁻¹, then mass of 2.5 mL solution in correct significant figures is :
 - (1) 53.75 g
- (2) 5375×10^{-3} g
- (3) 5.4 g
- (4) 5.38 g

Sol. Answer (3)

In case of multiplication and division, the final result should be reported as having the same number of significant digits as the number with least number of significant digits.

$$\therefore Density = \frac{Mass}{Volume}$$

So, Mass = 2.15×2.5

= 5.375

≈ 5.4 g

53. Given below are two statements : one is labelled as Assertion **(A)** and the other is labelled as Reason **(R)**.

Assertion (A):

Chlorine is an electron withdrawing group but it is ortho, para directing in electrophilic aromatic substitution.

Reason (R):

Inductive effect of chlorine destabilises the intermediate carbocation formed during the electrophilic substitution, however due to the more pronounced resonance effect, the halogen stabilises the carbocation at ortho and para positions.

In the light of the above statements, choose the most appropriate answer from the options given below :

- (1) (A) is not correct but (R) is correct.
- (2) Both (A) and (R) are correct and (R) is the correct explanation of (A).
- (3) Both (A) and (R) are correct but (R) is not the correct explanation of (A).
- (4) (A) is correct but (R) is not correct.

Sol. Answer (4)

Halogen are electron withdrawing groups due to high electronegativity. They have high electron withdrawing inductive effect. In electrophilic substitution reaction it shows both electron withdrawing inductive effect and electron donating resonance effect, but inductive effect overpowers resonance effect so, it deactivates benzene ring and directs the incoming electrophile to ortho and para positions. So, here assertion is correct and Reason is false.

54. Two half cell reactions are given below.

$$\text{Co}^{3+} + \text{e}^{-} \rightarrow \text{Co}^{2+}$$
 , $\text{E}^{\circ}_{\text{Co}^{2+}/\text{Co}^{3+}} = -1.81 \text{ V}$
 $2\text{Al}^{3+} + 6 \text{ e}^{-} \rightarrow 2\text{Al(s)}$, $\text{E}^{\circ}_{\text{Al/Al}^{3+}} = +1.66 \text{ V}$

The standard EMF of a cell with feasible redox reaction will be :

- (1) 3.47 V
- (2) + 7.09 V
- (3) + 0.15 V
- (4) + 3.47 V

Sol. Answer (4)

$$\mathsf{E}_{\mathsf{Cell}}^{\mathsf{0}} = \left(\mathsf{E}_{\mathsf{c}}^{\mathsf{0}} - \mathsf{E}_{\mathsf{a}}^{\mathsf{0}}\right)_{\mathsf{RP}}$$

- = 1.81 (-1.66)
- = 1.81 + 1.66
- = 3.47 V
- 55. Match List I with List II:

List - I

List - II

(Compounds) (Molecular formula)

- (a) Borax
- (i) NaBO₂
- (b) Kernite
- (ii) Na₂B₄O₇·4 H₂O
- (c) Orthoboric acid
- (iii) H₃BO₃
- (d) Borax bead
- (iv) Na₂B₄O₇·10 H₂O

Choose the **correct answer** from the options given below:

- (1) (a) (i), (b) (iii), (c) (iv), (d) (ii)
- (2) (a) (iv), (b) (ii), (c) (iii), (d) (i)
- (3) (a) (ii), (b) (iv), (c) (iii), (d) (i)
- (4) (a) (iii), (b) (i), (c) (iv), (d) (ii)

Sol. Answer (2)

Borax \rightarrow Na₂B₄O₇·10H₂O

Kernite \rightarrow Na₂B₄O₇·4H₂O

Orthoboric acid → H₃BO₃

Borax bead → NaBO₂

- **56.** The correct order of first ionization enthalpy for the given four elements is :
 - (1) C < F < N < O
- (2) C < N < F < O
- (3) C < N < O < F
- (4) C < O < N < F

Sol. Answer (4)

$$C \rightarrow 1s^2$$
, $2s^2 2p^2$

 $N \rightarrow 1s^2$, $2s^2 2p^3$ (more stable EC)

 $O \rightarrow 1s^2, 2s^2 2p^4$

 $F \to 1s^2, 2s^2 2p^5$

∴ order of first IE is C < O < N < F

57. Match List - I with List - II:

List - I

List - II

(Defects)

(shown by)

- (a) Frenkel defect
- (i) non-ionic solids and density of the solid decreases
- (b) Schottky defect
- (ii) non-ionic solids and density of the solid increases
- (c) Vacancy defect
- (iii) ionic solids and density of the solid decreases
- (d) Interstitial defect
- (iv) ionic solids and density of the solid

remains constant

Choose the **correct answer** from the options given below :

- (1) (a) (iv), (b) (iii), (c) (i), (d) (ii)
- (2) (a) (i), (b) (ii), (c) (iii), (d) (iv)
- (3) (a) (i), (b) (iii), (c) (ii), (d) (iv)
- (4) (a) (iv), (b) (iii), (c) (ii), (d) (i)

Sol. Answer (1)

Schottky and Frenkel defects are shown by ionic solids and in this density decreases and remains same respectively.

Vacancy and Interstitial defects are shown by non-ionic solid and in this density decreases and increases respectively.

Hence (a) - (iv), (b) - (iii), (c) - (i), (d) - (ii)

- **58.** Predict the order of reactivity of the following four isomers towards S_N2 reaction.
 - (I) CH₃CH₂CH₂CH₂CI
 - (II) CH₃CH₂CH(CI)CH₃
 - (III) (CH₃)₂CHCH₂CI
 - (IV)(CH₃)₃CCI
 - (1) (IV) > (II) > (III) > (I)
 - (2) (IV) > (III) > (II) > (I)
 - (3) (I) > (II) > (III) > (IV)
 - (4) (I) > (III) > (II) > (IV)
- Sol. Answer (4)

Greater the steric hindrance, lesser will be the rate of S_N2 reaction. So order of reactivity of S_N2 reaction is- (I) > (III) > (IV)

59. Match List - I with List - II:

List - I List - II (molecules) (shape)

- (a) NH₃
- (i) square pyramidal
- (b) CIF₃
- (ii) trigonal bipyramidal
- (iii) trigonal pyramidal
- (c) PCl₅(d) BrF₅
- (iv) T-shape

Choose the **correct answer** from the options given below:

- (2) (a) (ii), (b) (iii), (c) (iv), (d) (i)
- (3) (a) (iii), (b) (iv), (c) (ii), (d) (i)
- (4) (a) (iv), (b) (iii), (c) (i), (d) (ii)
- Sol. Answer (3)

 $NH_3 \rightarrow Trigonal pyramidal$

 $CIF_3 \rightarrow T$ -shape

PCl₅ → Trigonal bipyramidal

BrF₅ → Square pyramidal

- Which of the following reactions is a part of the large scale industrial preparation of nitric acid
 - (1) $Cu(NO_3)_2 + 2 NO_2 + 2 H_2O$

$$\xrightarrow{\text{Pt}}$$
 4HNO₃ + Cu

(2) NaNO₃ + H₂SO₄

$$\xrightarrow{\text{Pt}}$$
 NaHSO₄ + HNO₃

(3) 4 NH₃ + 5 O₂ (from air)

$$\xrightarrow{\text{Pt}}$$
 4 NO + 6 H₂O

(4) $4 \text{ HPO}_3 + 2 \text{ N}_2\text{O}_5$

$$\xrightarrow{\text{Pt}}$$
 4 HNO₃ + P₄O₁₀

Sol. Answer (3)

On large scale, nitric acid is prepared by Ostwald's process.

(i)
$$4NH_{3(g)} + 5O_2 \xrightarrow{Pt/Rh \text{ gauge} \atop 500K,9 \text{ bar}} 4NO_{(g)} + 6H_2O_{(g)}$$

- (ii) $2NO_{(q)} + O_{2(q)} \rightarrow 2NO_{2(q)}$
- (iii) $3NO_{2(a)} + H_2O_{(\ell)} \rightarrow 2HNO_{3(a0)} + NO$
- 61. Match List I with List II:

List - I List - II

- (a) Sodium (i) Toilet soap laurylsulphate
- (b) Cetyltrimethyl (ii) Non-ionic ammonium detergent chloride
- (c) Sodium stearate (iii) Anionic detergent
- (d) Polyethyleneglycyl (iv) Cationic stearate detergent

Choose the **correct answer** from the options give below:

- (1) (a) (iii), (b) (i), (c) (ii), (d) (iv)
- (2) (a) (iv), (b) (iii), (c) (i), (d) (ii)
- (3) (a) (i), (b) (iv), (c) (ii), (d) (iii)
- (4) (a) (iii), (b) (iv), (c) (i), (d) (ii)

Sol. Answer (4)

- (a) Sodium laurylsulphate → Anionic detergent
- (b) Cetyltrimethylammonium chloride → cationic detergent
- (c) Sodium stearate → Toilet soap
- (d) Polyethyleneglycyl stearate → Non-ionic detergent
- **62.** Which among the following is a thermoplastic polymer?
 - (1) Melamine polymer
 - (2) Bakelite
 - (3) Polythene
 - (4) Urea-formaldehyde resin

Sol. Answer (3)

Polythene, Polystyrene, polyvinyls etc. are thermoplastic polymers

63. $Na_2B_4O_7 \xrightarrow{heat} X + NaBO_2$

in the above reaction the product "X" is:

- (1) NaB₃O₅
- (2) H₃BO₃
- (3) B₂O₃
- (4) Na₂B₂O₅

Sol. Answer (3)

Borax on strong heating produces

Boric anhydride and sodium metaborate

 $Na_2B_4O_7 \xrightarrow{heat} B_2O_3 + 2NaBO_2$

64. One mole of an ideal gas at 300 K is expanded isothermally from 1 L to 10 L volume. ΔU for this process is :

(Use $R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1}$)

- (1) 0 J
- (2) 1260 J
- (3) 2520 J
- (4) 5040 J

Sol. Answer (1)

In isothermal expansion/compression of ideal gas

 $\Delta U = nC_v \Delta T$, $(\Delta T = 0)$

∴ ∆U = 0

65. Match List - I with List - II:

List - I List - II

(Complexes) (Types)

- (a) $[Co(NH_3)_5NO_2]Cl_2$ (i) ionisation and $[Co(NH_3)_5ONO]Cl_2$ isomerism
- (b) $[Cr(NH_3)_6][Co(CN)_6]$ (ii) coordination and $[Cr(CN)_6][Co(NH_3)_6]$ isomerism
- (c) $[Co(NH_3)_5(SO_4)]Br$ (iii) linkage and $[Co(NH_3)_5Br]SO_4$ isomerism
- (d) $[Cr(H_2O)_6]Cl_3$ and (iv) solvate $[Cr(H_2O)_5Cl]Cl_2 \cdot H_2O$ isomerism

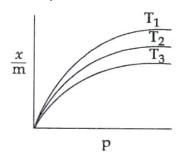
Choose the **correct answer** from the options given below:

- (1) (a) (iv), (b) (iii), (c) (ii), (d) (i)
- (2) (a) (iii), (b) (i), (c) (ii), (d) (iv)
- (3) (a) (ii), (b) (iii), (c) (iv), (d) (i)
- (4) (a) (iii), (b) (ii), (c) (i), (d) (iv)

Sol. Answer (4)

- (a) $[Co(NH_3)_5NO_2]Cl_2$ (iii) Linkage and $[Co(NH_3)_5ONO]Cl_2$ isomerism due to ambidentate ligand
- (b) $[Cr(NH_3)_6][Co(CN)_6]$ (ii) coordination and $[Cr(CN)_6][Co(NH_3)_6]$ isomerism due to exchange of ligands between coordination spheres
- (c) $[Co(NH_3)_5(SO_4)]Br$ (i) ionisation and $[Co(NH_3)_5Br]SO_4$ isomerism due to formation of different ions on
- (d) $[Cr(H_2O)_6]CI_3$ and $[Cr(H_2O)_5CI]CI_2 \cdot H_2O$
- ionisation

 (iv) solvate
 isomerism as
 no. of water
 molecules as
 ligand and
 water of
 crystalisation is
 different
- **66.** Shown below are adsorption isotherms for a gas 'X' at temperatures T_1 , T_2 and T_3 :



p and $\frac{x}{m}$ represent pressure and extent of adsorption, respectively. The correct order of temperatures for the given, adsorption is :

- (1) $T_1 = T_2 > T_3$
- (2) $T_1 > T_2 > T_3$
- (3) $T_3 > T_2 > T_1$
- (4) $T_1 = T_2 = T_3$

Sol. Answer (3)

As the temperature increases the extent of adsorption of a gas on solid surfaces decreases.

$$(T_3 > T_2 > T_1)$$

- **67.** 0.01 M acetic acid solution is 1% ionised, then pH of this acetic acid solution is :
 - (1) 1
- (2) 3
- (3) 2
- (4) 4
- Sol. Answer (4)

 $[H^+] = C.\alpha$

$$=0.01\times\frac{1}{100}=10^{-4}$$

$$pH = -log_{10}[H^+]$$

= 4

- **68.** The half life of a first order reaction is 2000 years. If the concentration after 8000 years is 0.02 M, then the initial concentration was:
 - (1) 0.04 M
- (2) 0.16 M
- (3) 0.32 M
- (4) 0.08 M
- Sol. Answer (3)

$$n = \frac{t}{t_{1/2}} = \frac{8000}{2000} = 4$$
, (no. of half lives)

$$\frac{[A_0]}{[A_t]} = 2^n, [A_0] = 0.02 \times 2^4 = 0.32 \text{ M}$$

69. The product formed from the following reaction sequence is :

$$\begin{array}{c|c} \text{NH}_2 & \text{(i)} & \text{(CH}_3\text{CO)}_2\text{O, pyridine} \\ \hline & \text{(ii)} & \text{LiAlH}_4 \\ \hline & \text{(iii)} & \text{H}_2\text{O} \\ \end{array}$$

Sol. Answer (2)

$$\begin{array}{c|c} NH_2 & NHCOCH_3 & NH & CH_3 \\ \hline \\ \hline \\ \hline \\ \hline \\ \hline \\ \hline \\ CH_3CO)_2O & \hline \\ \hline \\ \hline \\ \hline \\ \hline \\ \\ \hline \\ \\ \hline \\ \\ \hline \\ \\ \\ \end{array}$$

- **70** The decreasing order of boiling points of the following alkanes is:
 - (a) heptane
 - (b) butane
 - (c) 2-methylbutane
 - (d) 2-methylpropane
 - (e) hexane

Choose the correct answer from the options given below:

- (1) (a) > (e) > (c) > (b) > (d)
- (2) (a) > (c) > (e) > (d) > (b)
- (3) (c) > (d) > (a) > (e) > (b)
- (4) (a) > (e) > (b) > (c) > (d)
- Sol. Answer (1)

Boiling point of alkanes α molar mass.

Straight chain alkanes have more boiling point than branched alkanes.

Heptane has high molar mass and 2-methylpropane has low molar mass and is branched.

- **71.** The element used for welding metals with high melting points is :
 - (1) He
- (2) Cl₂
- (3) H_2
- (4) Ne
- Sol. Answer (3)

The oxy-hydrogen flame can produce the temperature of 4000 K. So atomic hydrogen is used for welding of metals with high melting point.

- 72. Decrease in size from left to right in actinoid series is greater and gradual than that in lanthanoid series due to:
 - (1) 5 f orbitals have greater shielding effect
 - (2) 4 f orbitals are penultimate
 - (3) 4 f orbitals have greater shielding effect
 - (4) 5 f orbitals have poor shielding effect
- Sol. Answer (4)

Due to large size of 5f orbitals their shielding effect is poor.

- 73. Which of the following reactions is not an example for nucleophilic addition - elimination reaction?
 - (1) $CH_3CHO + NH_3 \rightleftharpoons CH_3CH = NH + H_2O$
 - (2) CH₃CHO + NaHSO₃

$$\begin{array}{c} \text{OH} \\ \mid \\ \text{CH}_3 - \text{C} - \text{OSO}_2 \text{Na} \\ \mid \\ \text{H} \end{array}$$

(3) CH₃CHO + NH₂OH

$$\rightleftharpoons$$
 CH₃CH = N - OH + H₂O

(4) CH₃CHO + C₆H₅NHNH₂

$$\rightleftharpoons$$
 CH₃CH = N-NHC₆H₅ + H₂O

Sol. Answer (2)

In nucleophilic addition-elimination reactions along with the product water molecule is eliminated.

But in reaction of CH₃CHO and NaHSO₃ only addition takes place.

- **74.** CaCl₂ and Ca(OCl)₂ are components of :
 - (1) lime water
- (2) gypsum
- (3) Portland cement (4) bleaching powder
- Sol. Answer (4)

CaCl₂ and Ca(OCl)₂ are components of bleaching powder.

75. The product formed from the following reaction sequence is:

Sol. Answer (3)

$$\begin{array}{c|c}
O \\
C \\
H
\end{array}$$

$$\begin{array}{c|c}
OH \\
C \\
C \\
-NH_3
\end{array}$$

$$\begin{array}{c} OH \\ C \\ C \\ COOH \end{array} \xrightarrow{\text{(iii) NaOH,CaO}} \begin{array}{c} OH \\ C \\ H \end{array}$$

(NaOH + CaO) Sodalime is a decarboxylating reagent.

- 76. Flourine is a stronger oxidising agent than chlorine because:
 - (a) F-F bond enthalpy of has low dissociation.
 - (b) Flouride ion (F-) has high hydration enthalpy.
 - (c) Electron gain enthalpy of flourine is less negative than chlorine.
 - (d) Flourine has a very small size.

Choose the most appropriate answer from the options given:

- (1) (b) and (c) only
- (2) (a) and (b) only
- (3) (a) and (c) only
- (4) (a) and (d) only

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Sol. Answer (2)

$$X_{2(g)} \xrightarrow{\frac{\Delta H_{diss}}{\text{endothermic}}} 2X_{(g)} \xrightarrow{\frac{e^{-}}{\text{electron}}} X_{(g)}^{-} \xrightarrow{\frac{H_2O}{\Delta H}} X_{ad}^{-}$$

$$\underset{(exothermic)}{\underbrace{\frac{\Delta H_{diss}}{\Delta H}}} X_{ad}^{-}$$

By adding these values more energy is released for fluorine due to low bond dissociation enthalpy and high hydration enthalpy.

77. K_H value for some gases at the same temperature 'T' are given:

gas	K _H /k bar
Ar	40.3
CO ₂	1.67
НСНО	1.83 × 10 ⁻⁵
CH ₄	0.413

where K_H is Henry's Law constant in water. The order of their solubility in water is :

- (1) $HCHO < CH_4 < CO_2 < Ar$
- (2) $Ar < CO_2 < CH_4 < HCHO$
- (3) $Ar < CO_2 < CH_4 < HCHO$
- (4) $HCHO < CO_2 < CH_4 < Ar$

Sol. Answer (2)

Solubility of a gas
$$\propto \frac{1}{K_{H}}$$
 value

- **78.** Which of the following reactions is a decomposition redox reaction?
 - (1) $P_4(s) + 3OH^-(aq) + 3H_2O(l)$

$$\rightarrow$$
 PH₃(g) + 3H₂PO₂⁻ (aq)

(2) $2 \text{ Pb}(NO_3)_2(s)$

$$\rightarrow$$
 2PbO(s) + 4 NO₂(g) + O₂(g)

- (3) $N_2(g) + O_2(g) \rightarrow 2NO(g)$
- (4) $Cl_2(g) + 2OH^-(aq)$

$$\rightarrow$$
 CIO⁻(aq) + CI⁻(aq) + 4H₂O(I)

Sol. Answer (2)

Lead nitrate decomposed to give PbO, NO_2 and O_2 . In this Nitrogen atom oxidation state changes from +5 to +4 and oxygen changes from -2 to zero.

79. What is the hybridization shown by C₁ and C₂ carbons, respectively in the given compound?

(1) sp^3 and sp^3

(2) sp² and sp³

- (3) sp² and sp²
- (4) sp³ and sp²
- Sol. Answer (2)

$$OHC - CH = CH - CH_2 - COOCH_3$$

Ester group has more priority than aldehyde. So numbering should be done from left to right. C₁ has double bond and is sp² hybridised.

80. Match the reagents (**List - I**) with the product (**List - II**) obtained from phenol.

List-I List - II

- (a) (i) NaOH (ii) CO₂ (i) Benzoquinone (iii) H⁺
- (b) (i) Aqueous NaOH (ii) Benzene + CHCl₃ (ii) H⁺
- (c) Zn dust, Δ (iii) Salicyl aldehyde
- (d) Na₂Cr₂O₇, H₂SO₄ (iv) Salicylic acid

Choose the **correct answer** from the options given below:

- (1) (a) (iv), (b) (ii), (c) (i), (d) (iii)
- (2) (a) (iii), (b) (iv), (c) (i), (d) (ii)
- (3) (a) (ii), (b) (i), (c) (iv), (d) (iii)
- (4) (a) (iv), (b) (iii), (c) (ii), (d) (i)
- Sol. Answer (4)

- **81.** The correct order of bond angles in the following compounds/ species is :
 - (1) $CO_2 < NH_3 < H_2O < NH_4$
 - (2) $H_2O < NH_3 < NH_4 < CO_2$
 - (3) $H_2O < NH_4 < NH_3 < CO_2$
 - (4) $H_2O < NH_4 = NH_3 < CO_2$

Sol. Answer (2)

$$CO_2 \rightarrow 180^\circ$$

$$NH_4 \rightarrow 109.5^{\circ}$$

 $\ddot{N}H_3 \rightarrow 107^{\circ}$ (N atom has lone pair)

 $H_2\ddot{O}$: \rightarrow 104.5° (oxygen atom has two lone pairs)

Due to lone pair-lone pair repulsions the bond angle in water decreases more.

82. Match List - I with List - II:

List - I List - II

(Reaction) (Product formed)

(i) Benzaldehyde

- (a) Gabriel synthesis
- (ii) Ethers
- (b) Kolbe synthesis

(c) Williamson synthesis

- (iii) Primary amines
- (d) Etard reaction
- (iv) Salicylic acid

Choose the correct answer from the options given below

- (1) (a) (iii), (b) (iv), (c) (ii), (d) (i)
- (2) (a) (iii), (b) (i), (c) (ii), (d) (iv)
- (3) (a) (ii), (b) (iii), (c) (i), (d) (iv)
- (4) (a) (iv), (b) (iii), (c) (i), (d) (ii)

Sol. Answer (1)

(a) Gabriel synthesis

(b) Kolbe synthesis

$$\begin{array}{ccc}
OH & ONa & OH \\
O & \stackrel{\text{NaOH}}{\longrightarrow} & O & OH \\
O & \stackrel{(i) CO_2}{\longrightarrow} & OH \\
O & OH \\
Salicylic acid
\end{array}$$

(c) Williamson synthesis

$$R \longrightarrow X + R'ONa \longrightarrow R'OR + NaX$$

(d) Etard reaction

$$\bigcirc \overset{\mathsf{CH}_3}{\longrightarrow} + \mathsf{CrO}_2\mathsf{Cl}_2 \xrightarrow{\mathsf{CS}_2} \bigcirc \overset{\mathsf{CH}_3\mathsf{O}^+}{\longrightarrow} \bigcirc \overset{\mathsf{H}_3\mathsf{O}^+}{\longrightarrow} \bigcirc \overset{\mathsf{CHO}}{\longrightarrow} \overset{\mathsf{$$

83. If first ionization enthalpies of elements X and Y are 419 kJ mol⁻¹ and 590 kJ mol⁻¹, respectively and second ionization enthalpies of X and Y are 3069 kJ mol⁻¹ and 1145 kJ mol⁻¹, respectively.

Then correct statement is:

- (1) Both X and Y are alkaline earth metals.
- (2) X is an alkali metal and Y is an alkaline earth metal.
- (3) X is an alkaline earth metal and Y is an alkali metal.
- (4) Both X and Y are alkali metals.

Sol. Answer (2)

X-is alkali metal as it has large size, it's IE is less.

After loss of one electron, it gets inert gas configuration. So its IE₂ is very high.

Y is alkaline earth metal. It's IE is more than alkali metal due to stable ns² configuration. But its IE₂ is lower than alkali metal.

- **84.** The **incorrect** statement about denaturation of proteins is :
 - (1) Uncoiling of the helical structure takes place.
 - (2) It results due to change of temperature and/ or pH
 - (3) It results in loss of biological activity of proteins.
 - (4) A protein is formed from amino acids linked by peptide bonds.

Sol. Answer (4)

Protein formation is not related to the denaturation of proteins.

85. Four gas cylinders containing He, N₂, CO₂ and NH₃ gases separately are gradually cooled from a temperature of 500 K. Which gas will liquify first?

(Given T_c in K - He : 5.3, N_2 : 126, CO_2 : 304.1 and NH_3 : 405.5)

- (1) NH₃
- (2) He
- (3) N_2
- (4) CO₂

Sol. Answer (1)

Ease of Liquefaction ∞ Critical Temperature So, NH₃ will liquify first.

Chemistry: Section-B (Q. No. 86 to 100)

86. Given below are two statements: one is labelled as **Assertion (A)** and the other is labelled as **Reason (R)**.

Assertion (A) : The metal carbon bond in metal carbonyls possesses both σ and π character.

Reason (R): The ligand to metal bond is a π bond and metal to ligand bond is a σ bond.

In the Light of the above statements, choose the **most appropriate** answer from the options given below:

- (1) (A) is not correct but (R) is correct
- (2) Both (A) and (R) are correct and (R) is the correct explanation of (A)
- (3) Both (A) and (R) are correct but (R) is not the correct explanation of (A)
- (4) (A) is correct but (R) is not correct

Sol. Answer (4)

Metal-carbon bond in metal carbonyls possesses both σ and π character, So the assertion is correct.

The ligand to metal bond is σ bond and metal to ligand bond is π bond, So the reason is correct.

87. Match List - I with List - II:

List - I

List - II

- (a) Biochemical oxygen demand
- (i) oxidising mixture
- (b) Photochemical smog
- (ii) polar stratospheric cloud
- (c) Classical smog
- (iii) organic matter In water
- (d) Ozone layer depletion
- (iv) reducing mixture

Choose the **correct answer** from the options given below:

- (1) (a) (iv), (b) (iii), (c) (ii), (d) (i)
- (2) (a) (i), (b) (iv), (c) (ii), (d) (iii)
- (3) (a) (iii), (b) (iv), (c) (i), (d) (ii)
- (4) (a) (iii), (b) (i), (c) (iv), (d) (ii)

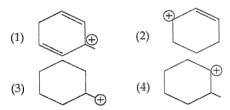
Sol. Answer (4)

Biochemical oxygen demand – Organic matter in water

Photochemical smog – Oxidising in nature Classical smog – Reducing in nature

Ozone layer depletion – Polar stratospheric cloud

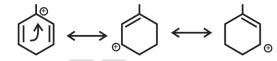
88. Which of the following is the most stable carbocation?



Sol. Answer (1)

Stability of carbocation ∞ No of α -H

∞ No of resonating structures



89. Given below are two statements:

Statement I: Cr²⁺ is oxidising and Mn³⁺ is reducing in nature.

Statement II: Sc³⁺ compounds are repelled by the applied magnetic field.

In the light of the above statements, choose the **most appropriate** answer from the options given below:

- Statement I is incorrect but Statement II is correct
- (2) Both Statement I and Statement II are correct
- (3) Both Statement I and Statement II are incorrect
- (4) Statement I is correct but Statement II is incorrect

Sol. Answer (1)

$$Cr^{+2}$$
: [Ar] $3d^4$

 Cr^{+2} is reducing as its configuration changes from d^4 to d^3 ($t_{2\alpha}^3$)

Mn⁺³ is oxidising in nature.

Mn⁺²: [Ar] 3d⁵ (extra stability)

Statement I is incorrect

Sc+3: [Ar]

diamagnetic - repelled by magnetic field.

Statement (II) is correct.

90. K_p for the following reaction is 3.0 at 1000 K.

$$CO_2(g) + C(s) \Longrightarrow 2CO(g)$$

What will be the value of K_c for the reaction at the same temperature ?

(Given $- R = 0.083 L bar K^{-1} mol^{-1}$)

- (1) 3.6
- (2) 0.36
- (3) 3.6×10^{-2}
- $(4) 3.6 \times 10^{-3}$
- Sol. Answer (3)

$$CO_2(g) + C(s) \implies 2CO(g)$$

$$K_P = K_c (RT)^{\Delta n_g} (\Delta n_g = 2 - 1)$$

 $3 = K_c (0.083 \times 1000)$

$$K_c = \frac{3}{0.083 \times 1000} = 3.6 \times 10^{-2}$$

91. A vessel contains 3.2 g of dioxygen gas at STP (273.15 K and 1 atm pressure). The gas is now transferred to another vessel at constant temperature, where pressure becomes one third of the original pressure. The volume of new vessel in L is:

(Given - molar volume at STP is 22.4 L)

- (1) 67.2
- (2) 6.72
- (3) 2.24
- (4) 22.4
- Sol. Answer (2)

Moles of oxygen =
$$\frac{3.2}{32}$$
 = 10⁻¹ mole

Volume at STP = $10^{-1} \times 22.4 = 2.24 L$

 $P_1 = 1$ atm $V_1 = 2.24$ L

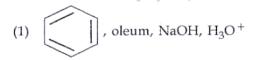
$$P_2 = \frac{1}{3}$$
 atm $V_2 = ?$

 $P_1V_1 = P_2V_2$

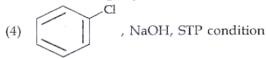
$$1 \times 2.24 L = \frac{1}{3} \times V_2$$

 $V_2 = 3 \times 2.24 L = 6.72 L$

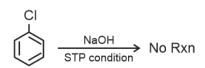
92. Which one of the following reaction sequence is **incorrect** method to prepare phenol?

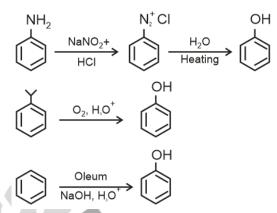


- (2) Aniline, NaNO2+HCl, H2O, heating
- (3) Cumene, O₂, H₃O⁺



Sol. Answer (4)





93. For a chemical reaction

$$4A + 3B \rightarrow 6C + 9D$$

rate of formation of C is 6×10^{-2} mol L⁻¹ s⁻¹ and rate of disappearance of A is 4×10^{-2} mol L⁻¹ s⁻¹. The rate of reaction and amount of B consumed in interval of 10 seconds, respectively will be :

- (1) $10 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1}$ and $30 \times 10^{-2} \text{ mol L}^{-1}$
- (2) $1 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1}$ and $30 \times 10^{-2} \text{ mol L}^{-1}$
- (3) $10 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1}$ and $10 \times 10^{-2} \text{ mol L}^{-1}$
- (4) $1 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1}$ and $10 \times 10^{-2} \text{ mol L}^{-1}$
- Sol. Answer (2)

$$4A + 3B \rightarrow 6C + 9D$$

$$r=-\frac{1}{4}\,\frac{d[A]}{dt}=+\frac{1}{6}\,\frac{d[C]}{dt}$$

$$r = \frac{1}{6} \times 6 \times 10^{-2} = 1 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1}$$

$$r = -\frac{1}{3} \frac{d[B]}{dt}$$

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$$\frac{-d[B]}{dt} = 3 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1}$$

B consumed in 10 sec = $3 \times 10^{-2} \times 10$

$$= 30 \times 10^{-2} \text{ mol L}^{-1}$$

94. Standard electrode potential for the cell with cell reaction

$$Zn(s) + Cu^{2+}(aq) \rightarrow Zn^{2+}(aq) + Cu(s)$$

is 1.1 V. Calculate the standard Gibbs energy change for the cell reaction.

(Given $F = 96487 \text{ C mol}^{-1}$)

- (1) -200.27 J mol⁻¹ (2) -200.3
 - (2) -200.27 kJ mol⁻¹
- (3) $-212.27 \text{ kJ mol}^{-1}$ (4) $-212.27 \text{ J mol}^{-1}$
- Sol. Answer (3)

$$Zn(s) + Cu^{+2}(aq) \rightarrow Zn^{+2}(aq) + Cu(s) v$$

$$E_{cell}^{\circ} = 1.1 \text{ V}$$

$$n = 2$$

$$\Delta G^{\circ} = -nFE_{cell}^{\circ}$$

$$= -2 \times 96487 \times 1.1$$

$$= -212.27 \text{ kJ}$$

95. The products A and B in the following reaction sequence are :

Ph
$$\xrightarrow{\text{(i) } \text{HBr}} A \xrightarrow{\text{(i) } \text{SOCl}_2} B$$

 $\xrightarrow{\text{(ii) } \text{Mg, dry ether}} A \xrightarrow{\text{(ii) } \text{CH}_3\text{NH}_2} B$

(1)
$$A = \bigcirc OH$$
 ; $B = \bigcirc OH$;

(2)
$$A = \bigcirc OH$$
; OH ; OH

(3)
$$A = \bigcup_{O} OH$$
; $H = \bigcup_{O} CH_3$

(4)
$$A = \begin{pmatrix} OH \\ H \\ N \end{pmatrix}$$

Sol. Answer (3)

$$\begin{array}{c} \text{Br} \\ \text{Mg} \\ \text{dry ether} \\ \text{O=C=O} \\ \text{OO} \\ \text{CO}_2, \text{H}_3\text{O} \\ \text{OO} \\ \text{OH} \\$$

96. Which one of the following is not a calcination reaction?

(1)
$$CaCO_3 + 2HCI \xrightarrow{\Delta} CaCl_2 + H_2O + CO_2$$

(2)
$$ZnCO_3 \xrightarrow{\Delta} ZnO + CO_2$$

(3)
$$\operatorname{Fe_2O_3} \cdot \operatorname{xH_2O} \xrightarrow{\Delta} \operatorname{Fe_2O_3} + \operatorname{xH_2O}$$

(4)
$$CaCO_3 \cdot MgCO_3 \xrightarrow{\Delta} CaO + MgO + 2CO_2$$

Sol. Answer (1)

Calcination involves heating in absence of air and the volatile matter escaped leaving behind the metal oxide

$$ZnCO_{3(s)} \xrightarrow{\Delta} ZnO_{(s)} + CO_{2(g)}$$

$$Fe_2O_3 \cdot XH_2O \xrightarrow{\Delta} Fe_2O_{3(s)} + XH_2O_{(g)}$$

$$CaCO_3 \cdot MgCO_3 \xrightarrow{\Delta} CaO_{(s)} + MgO_{(s)} + 2CO_{2(g)}$$

- **97.** The **incorrect** method for the synthesis of alkenes is :
 - (1) treating vicinal dihalides with Zn metal
 - (2) treatment of alkynes with Na in liquid NH3
 - (3) heating alkyl halides with alcoholic KOH
 - (4) treating alkyl halides in aqueous KOH solution

Sol. Answer (4)

(4)
$$R - CI \xrightarrow{aq KOH} R - OH$$

(3)
$$CH_3-CH_2-CH-CH_3 \xrightarrow{alc} CH_3-CH = CH-CH_3 + CH_3-CH_2 -CH=CH_2$$

(2)
$$R-C \equiv C-R \xrightarrow{\text{Na}} \xrightarrow{\text{Liq NH}_3} \xrightarrow{R} C=C \xrightarrow{H}$$

(1)
$$CH_2-CH_2 \xrightarrow{X} CH_2=CH_2$$
 $CH_2=CH_2$

98. When electromagnetic radiation of wavelength 300 nm falls on the surface of a metal, electrons are emitted with the kinetic energy of 1.68 × 10⁵ J mol⁻¹. What is the minimum energy needed to remove an electron from the metal?

(h = $6.626 \times 10^{-34} \text{ Js}, c = 3 \times 10^8 \text{ ms}^{-1},$

 $N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$

- (1) $2.31 \times 10^5 \text{ J mo}1^{-1}$
- (2) $2.31 \times 10^6 \text{ J mo}1^{-1}$
- (3) $3.84 \times 10^4 \text{ J mo}1^{-1}$
- (4) $3.84 \times 10^{-19} \,\mathrm{J \ mo1^{-1}}$
- Sol. Answer (1)

$$E_p = \phi + K.E$$

 $E_P - K.E = \phi$

$$E_P = \frac{hc}{\lambda} = 3.975 \times 10^5 \text{ J mol}^{-1}$$

 $K.E = 1.68 \times 10^5 \text{ J mol}^{-1}$

 $\phi = (3.975 - 1.68) \times 10^5$

= $2.295 \times 10^5 \approx 2.31 \times 10^5 \text{ J mol}^{-1}$

- **99.** What fraction of Fe exists as Fe(III) in Fe_{0.96}O? (Consider Fe_{0.96}O to be made up of Fe(II) and Fe(III) only)
 - (1) $\frac{1}{20}$
- (2) $\frac{1}{12}$
- (3) 0.08
- (4) $\frac{1}{16}$
- Sol. Answer (2)

 $Fe_{0.96}O = Fe_{96}O_{100}$

Let us consider Fe in $Fe^{+2} = x$

Fe in Fe⁺³ = (96 - x)

Total +ve charge = total -ve charge

$$(96 - x) \times 3 + 2x = 200$$

$$288 - 3x + 2x = 200$$

x = 88

$$Fe^{+3} = 96 - 88 = 8$$

Fraction of Fe⁺³ = $\frac{8}{96} = \frac{1}{12}$

100. The incorrect method to synthesize benzaldehyde is :

followed by H₃O+

(3)
$$OC_2H_5$$
, DIBAL-H,

followed by H2O

followed by H₃O⁺ in CS₂

Sol. Answer (1)

$$(2) \qquad C \qquad CI \qquad \xrightarrow{H_2, Pd \text{ BaSO}_4} \qquad C \qquad H$$

Rosenmund Reaction

$$(4) \bigcirc CH_3 \xrightarrow{CrO_2Cl_2} \bigcirc H$$

Etard reaction