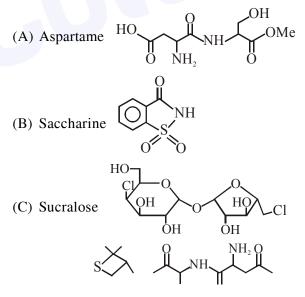


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| 3 | | | |
|------|--|--|--|
| 4. | A chemist has 4 samples of artificial sweetener | 5. 'X' melts at low temperature and is a bad | |
| | A, B, C and D. To identify these samples, he | conductor of electricity in both liquid and solid | |
| | performed certain experiments and noted the | state. X is : | |
| | following observations : | (1) Carbon tetrachloride (2) Mercury | |
| | (i) A and D both form blue-violet colour with | (3) Silicon carbide (4) Zinc sulphide | |
| | ninhydrin. | NTA Ans. (1) | |
| | (ii) Lassaigne extract of C gives positive | Sol. CCl_4 is molecular solid so does not conduct | |
| | AgNO ₃ test and negative $Fe_4[Fe(CN)_6]_3$ test. | electricity in liquid & solid state. | |
| | (iii)Lassaigne extract of B and D gives positive | 6. According to the following diagram, A reduces | |
| | sodium nitroprusside test | BO_2 when the temperature is : | |
| | Based on these observations which option is | | |
| | correct ? | 600 - | |
| | (1) A : Aspartame ; B : Saccharin ; | $ \begin{array}{c} -600 \\ \hline \\ 0 \\ 0 \\ 0 \\ 0 \\ 0 \\ 0 \\ 0 \\ 0 \\$ | |
| | C : Sucralose ; D ; Alitame | $\overset{\swarrow}{O}_{O}$ -1000 A + O ₂ \rightarrow AO ₂ | |
| | (2) A : Alitame ; B : Saccharin ; | | |
| | C : Aspartame ; D ; Sucralose | $-1200 - B + O_2 \rightarrow BO_2$ | |
| | (3) A : Saccharin ; B : Alitame ; | | |
| | C : Sucralose ; D ; Aspartame | $\begin{array}{cccccccccccccccccccccccccccccccccccc$ | |
| | (4) A : Aspartame ; B : Alitame ; | $(1) < 1400 \ ^{\circ}C$ | |
| | C : Saccharin ; D ; Sucralose | $(2) > 1400 \ ^{\circ}\mathrm{C}$ | |
| NTA | Ans. (1) | (3) < 1200 °C | |
| Sol. | (i) Blue voilet color with Ninhydrine \rightarrow amino | (4) > 1200 °C but < 1400 °C | |
| | acid derivative. So it cannot be saccharide or | NTA Ans. (2) | |

- sucralose. (ii) Lassaigne extract give +ve test with AgNO₃. So Cl is present, -ve test with $Fe_4[Fe(CN)_6]_3$ means N is absent. So it can't be Aspartame or Saccharine or
- (iii) Lassaigne solution of B and D given +ve sodium nitroprusside test, so it is having S, so it is Saccharine and Alitame.

Alitame, so C is sucralose.



Sol. A reduces BO_2 when temperature is above 1400°C because above 1400°C A has more ve ΔG° for AO₂ formation than B to BO₂ formation.

7. The K_{sp} for the following dissociation is 1.6×10^{-5}

$$PbCl_{2(s)} \rightleftharpoons Pb_{(aq)}^{2+} + 2Cl_{(aq)}^{-}$$

Which of the following choices is correct for a mixture of 300 mL 0.134 M Pb(NO₃)₂ and 100 mL 0.4 M NaCl ?

(1) $Q < K_{sp}$

(2)
$$Q > K_{sj}$$

(3)
$$Q = K_{s_1}$$

(4) Not enough data provided

NTA Ans. (2)

Sol.
$$\left[Pb^{2+} \right] = \frac{300 \times 0.134}{400} = 1.005 \times 10^{-1} \text{ M}$$

$$\left[\text{Cl}^{-} \right] = \frac{100 \times 0.4}{400} = 10^{-1} \text{ M}$$

$$PbCl_{2(s)} \Longrightarrow Pb_{(aq.)}^{+2} + 2Cl_{(aq.)}^{-}$$



| 0 | $\left[\mathbf{D} \left[\left(\mathbf{E} \right) \left(\mathbf{O} \right) \right] \right] = \left[\mathbf{D} \left[\left(\mathbf{E} \right) \left(\mathbf{O} \right) \right] \right] = \left[\mathbf{D} \left[\left(\mathbf{E} \right) \left(\mathbf{O} \right) \right] \right] = \left[\mathbf{D} \left[\left(\mathbf{E} \right) \left(\mathbf{O} \right) \right] \right] = \left[\mathbf{D} \left[\left(\mathbf{E} \right) \left(\mathbf{O} \right) \right] \right] = \left[\mathbf{D} \left[\mathbf{E} \right] \right] = \left[\mathbf{D} \left[\mathbf{E} \right] \right] = \left[\mathbf{D} \left[\mathbf{E} \right] \left(\mathbf{O} \right) \left(\mathbf{E} \right) \right] = \left[\mathbf{D} \left[\mathbf{E} \right] \right] = \left[\mathbf{D} \left[\mathbf{E} \right] \right] = \left[\mathbf{D} \left[\mathbf{E} \right] \left(\mathbf{E} \right) \left(\mathbf{E} \right) \right] = \left[\mathbf{D} \left[\mathbf{E} \right] \right] = \left[\mathbf{D} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \right] = \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \left[\mathbf{E} \right] \right] = \left[\mathbf{E} \left[\mathbf{E} \left$ | | |
|------|--|--|--|
| 8. | $[Pd(F)(Cl)(Br)(I)]^{2-}$ has n number of geometrical | | |
| | isomers. Then, the spin-only magnetic moment | | |
| | and crystal field stabilisation energy [CFSE] of | | |
| | $[Fe(CN)_6]^{n-6}$, respectively, are: | | |
| | [Note : Ignore the pairing energy] | | |
| | (1) 2.84 BM and $-1.6 \Delta_0$ | | |
| | (2) 1.73 BM and $-2.0 \Delta_0$ | | |
| | (3) 0 BM and $-2.4 \Delta_0$ | | |
| | (4) 5.92 BM and 0 | | |
| | Ans. (2) $(D = (D $ | | |
| Sol. | $[Pb(F)(Cl)(Br)(I)]^{2-}$ have three geometrical | | |
| | isomer so formula for $[Fe(CN)_6]$ | | |
| | is $[Fe(CN)_6]^{3-}$ and CFSE for this complex is | | |
| | $\mathrm{Fe}^{3\oplus} \Rightarrow 3\mathrm{d}^{5}4\mathrm{s}^{\circ}$ | | |
| | | | |
| | | | |
| | | | |
| | | | |
| | | | |
| | Magnetic Moment = $\sqrt{3}$ | | |
| | = 1.73 B.M | | |
| | $CFSE = [(-0.4 \times 5) + (0.6 \times 0)] \Delta_0$ | | |
| | $= -2.0 \Delta_0$ | | |
| 9. | If the magnetic moment of a dioxygen species | | |
| | is 1.73 B.M, it may be : | | |
| | (1) O_2^- or O_2^+ | | |
| | 2 2 | | |
| | (2) $O_2 \text{ or } O_2^+$ | | |
| | (3) O_2 or O_2^- | | |
| | (4) O_2 , O_2^- or O_2^+ | | |
| NTA | Ans. (1) | | |
| Sol. | number of magnetic moment | | |
| | unpaired electron | | |
| | O_2^{\ominus} 1 1.73 B.M | | |
| | O ₂ ⊕ 1 1.73 B.M | | |
| | O ₂ 2 2.83 BM | | |
| 10. | If enthalpy of atomisation for $Br_{2(1)}$ is x kJ/mol | | |
| | and bond enthalpy for Br_2 is y kJ/mol, the | | |
| | relation between them : | | |
| | (1) is $x = y$ (2) is $x < y$ | | |
| | (3) does not exist (4) is $x > y$ | | |
| NTA | Ans. (4) | | |
| | | | |

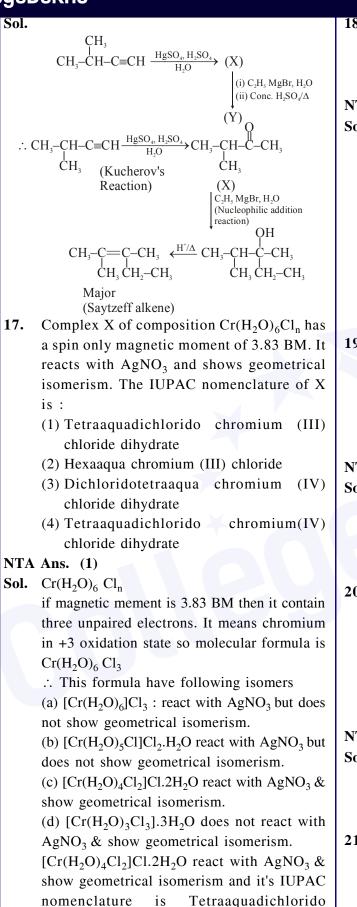
Sol. Enthalpy of atomisation of $Br_2(l)$ $\begin{array}{c} \operatorname{Br}_{2}(l) \xrightarrow{\Delta \operatorname{H}_{\operatorname{vap}}} \operatorname{Br}_{2}(g) \xrightarrow{\Delta \operatorname{H}_{\operatorname{BE}}} 2\operatorname{Br}(g) \\ \uparrow & & \uparrow \\ \Delta \operatorname{H}_{\operatorname{atom.}} \end{array}$ $\Delta H_{atom} = \Delta H_{vap} + \Delta H_{BE}$ $x = \Delta H_{vap} + y$ So, x > y11. The increasing order of basicity for the following intermediates is (from weak to strong) (i) $H_3C - C\Theta$ CH_3 $C\Theta$ CH_3 (ii) $H_2C = CH - \overset{\Theta}{C}H_2$ (iii) $HC \equiv \overset{\Theta}{C}$ (iv) $\overset{\Theta}{C}H_3$ (v) $\overset{\Theta}{C}N$ (1) (v) < (i) < (iv) < (ii) < (iii) (2) (iii) < (i) < (ii) < (iv) < (v) (3) (v) < (iii) < (ii) < (iv) < (i)(4) (iii) < (iv) < (ii) < (i) < (v) NTA Ans. (3) ÇH₃ **Sol.** $CH_3 - C_{13} - C_{13$ HC≡C⊖ (ii) (i) (iii) ⊖ C≡N CH_3^{\ominus} (iv) (v) Basic strength order : (i) > (iv) > (ii) > (iii) > (v)12. B has a smaller first ionization enthalpy than Be. Consider the following statements : (I) It is easier to remove 2p electron than 2s electron (II) 2p electron of B is more shielded from the nucleus by the inner core of electrons than the 2s electrons of Be. (III) 2s electron has more penetration power than 2p electron. (IV) atomic radius of B is more than Be (Atomic number B = 5, Be = 4) The correct statements are : (1) (I), (II) and (III)

(2) (II), (III) and (IV)

(3) (I), (III) and (IV) $% \left(IV\right) \left($

Sol. Be \Rightarrow 1s² 2s² Sol. $B \Rightarrow 1s^2 2s^2 2p^1$ NH B has a smaller size than Be NaNO₂/HCl it is easier to remove 2p electron than 2s 273_278K electron due to less pentration effect of 2p than (Diazotisation) (m-Bromo benzene 2s. diazonium chloride) 2p electron of Boron is more shielded from the nucleus by the inner core of electron than the HNO₂ Cu_2Br_2 Sand Mayer's (X)2s electron of Be Brreaction Br (Nitration) B has a smaller size than Be Which of these will produce the highest yield 15. 13. The acidic, basic and amphoteric oxides, in Friedel Crafts reaction? respectively, are : CONH₂ (1) MgO, Cl₂O, Al₂O₃ ີ) (3) ໂ (2)(2) Cl₂O, CaO, P₄O₁₀ (1)(3) Na₂O, SO₃, Al₂O₃ NTA Ans. (3) (4) N₂O₃, Li₂O, Al₂O₃ Sol. (Deactivated ring due to *.*.. NTA Ans. (4) -NH₂ -R effect of amide) Sol. 1. MgO Basic Cl₂O Acidic Al₂O₃ amphoteric (l.p.e. of $-NH_2$ and -QH2. Cl₂O Acidic CaO Basic group do acid-base reaction P₄O₁₀ Acidic with lewis acid AlCl₃, so ring 3. Na₂O Basic is deactivated) SO₃ Acidic Al₂O₃ amphoteric (Highest yield produced) 4. N₂O₃ Acidic Li₂O Basic The major product (Y) in the following 16. Al₂O₃ amphoteric reactions is : The major product Z obtained in the following 14. reaction scheme is : CH₃ $CH_3 - CH - C \equiv CH \xrightarrow{HgSO_4, H_2SO_4} X \rightarrow X$ (i) C_2H_5MgBr, H_2O (ii)Conc. H_2SO_4/Δ NH_2 (1) $H_3C - C - CH - CH_3$ C_2H_5 NaNO2/HCl $\rightarrow X - \frac{Cu_2Br_2}{Cu_2Br_2}$ Br ÇH3 (2) $CH_3 - CH - CH - CH_3$ O_2 ĊH₃ (1)(2) Br (3) $CH_3 - C = C - CH_3$ CH_2CH_3 $CH_3 - CH - C = CH_2$ (4) $CH_3 - CH - C = CH_2$ NO_2 (3) Br-(4)B





| 18. The compound that cannot act bot | h as oxidising |
|---|----------------|
|---|----------------|

- and reducing agent is :
- (1) H_2O_2 (2) H_2SO_3 (3) HNO_2 (4) H_3PO_4

NTA Ans. (4)

- Sol. (i) H_2O_2 act as oxidising agent as well as reducing agent depending on condition.
 - (ii) H_2SO_3 act as oxidising agent as well as reducing agent depending on condition.
 - (iii) HNO_2 act as oxidising agent as well as reducing agent depending on condition.
 - (iv) H_3PO_4 can not act both as oxidising and reducing agent.

H₃PO₄ can act as only oxidising agent.

 $H_3PO_4 \implies 3H^+ + PO_4^{3-}$

- **19.** The de Broglie wavelength of an electron in the 4th Bohr orbit is :
 - (1) $8\pi a_0$
 - (2) $2\pi a_0$
 - (3) $4\pi a_0$
 - (4) 6πa₀
- NTA Ans. (1)
- **Sol.** $2\pi r = n\lambda$
 - for $n = 1, r = a_0$ $n = 4, r = 16a_0$ So, $2\pi \times 16a_0 = 4 \times \lambda$

$$\lambda = 8\pi a_0$$

- 20. The electronic configurations of bivalent europium and trivalent cerium are
 - (atomic number : Xe = 54, Ce = 58, Eu = 63)
 - (1) [Xe] $4f^4$ and [Xe] $4f^9$
 - (2) [Xe] $4f^7$ and [Xe] $4f^1$
 - (3) [Xe] $4f^7\ 6s^2$ and [Xe] $4f^2\ 6s^2$
 - (4) [Xe] 4f² and [Xe] 4f⁷
- NTA Ans. (2)
- Sol. $\operatorname{Eu}_{63} \Rightarrow [\operatorname{Xe}] 4f^7 5d^\circ 6s^2$ $\operatorname{Eu}^{2\oplus} \Rightarrow [\operatorname{Xe}] 4f^7$ $\operatorname{Ce}_{58} \Rightarrow [\operatorname{Xe}] 4f^1 5d^1 6s^2$ $\operatorname{Ce}^{3\oplus} \Rightarrow [\operatorname{Xe}] 4f^1$
- 21. The hardness of a water sample containing 10^{-3} M MgSO₄ expressed as CaCO₃ equivalents (in ppm) is _____. (molar mass of MgSO₄ is 120.37 g/mol)

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Sol. 1 Litre has 10^{-3} moles MgSO₄ So , 1000 litre has 1 mole MgSO₄

 $= 1 \text{ mole CaCO}_3$

= 100 ppm

22. The molarity of HNO_3 in a sample which has density 1.4 g/mL and mass percentage of 63% is _____. (Molecular Weight of $HNO_3 = 63$)

NTA Ans. (14.00)

Sol. 100 gm soln \rightarrow 63 gm HNO₃

 $\frac{100}{1.4}\text{mL} \rightarrow 1 \text{ mole HNO}_3$

Molarity =
$$\frac{1}{\frac{100}{1.4} \times \frac{1}{1000}} = 14$$
M

- 23. 108 g of silver (molar mass 108 g mol⁻¹) is deposited at cathode from AgNO₃(aq) solution by a certain quantity of electricity. The volume (in L) of oxygen gas produced at 273 K and 1 bar pressure from water by the same quantity of electricity is _____.
- NTA Ans. (5.66 to 5.68)

Sol. gm eq. of Ag =
$$\frac{108}{108}$$
 =

gm eq. of $O_2(g) = 1$

Volume of $O_2(g) = 22.7 \times \frac{1}{4} = 5.675$ litre

24. The mass percentage of nitrogen in histamine is

NTA Ans. (37.80 to 38.20)

M.F. of Histamine is $C_5H_9N_3$ Molecular mass of Histamine is 111

 $/NH_2$

Now, mass % of nitrogen = $\left(\frac{42}{111}\right) \times 100$

- = 37.84%
- 25. How much amount of NaCl should be added to 600 g of water ($\rho = 1.00$ g/mL) to decrease the freezing point of water to -0.2 °C ? _____. (The freezing point depression constant for water = 2K kg mol⁻¹)
- NTA Ans. (1.74 to 1.76 or 0.03)
- **Sol.** $\Delta T_f = i \times m \times K_f$

$$0.2 = 2 \times 2 \times \frac{w/58.5}{600/1000}$$

w = 1.755 gm