R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

Chapter 8: Volumetric analysis Q1. **Sol:** Strength in g / lit = density = ? We have, NHCI = 3 $M_{HCI} = \frac{3}{1} = 3$ (::, v.f·=1) $N_{H_2SO_4} = \frac{1}{2}$ $\therefore M_{H_2SO_4} = \frac{N_{H_2SO_4}}{vf} = \frac{1/2}{2} = \frac{1}{4}$ wt. of HCl / lit = $3 \times Mol.$ wt = $3 \times 36.5 = 109.5$ g /lit Ans $\frac{\text{moles of H}_2\text{SO}_4}{\text{lit}} = \frac{1}{4} \implies \frac{\text{wt of H}_2\text{SO}_4/\text{lit}}{\text{Mol.wt H}_2\text{SO}_4} = \frac{1}{4}$ wt. of H₂SO₄ / lit = $\frac{1}{4} \times 98 = 24.5 \text{ g/lit}$ Ans Q2. Sol: NaOH + $H_2SO_4 \rightarrow Na_2SO_4 + H_2O$ 10 ml 1 M 1 M. meq. of NaOH = meq. of H_2SO_4 $10 \times 1 \times 1 = V \times 1 \times 2$ $V = \frac{10}{2} = 5 \,\text{mL}$ Ans Q3. Sol: V = 2 lit of NH₃, T = $13^{\circ}C = 286$ K, P = 0.9 atm moles = $\frac{PV}{RT} = \frac{0.9 \times 2}{0.821 \times 286} = 0.076$ mole $NH_3 \longrightarrow H_2SO_4 \longrightarrow (NH_4)_2SO_4$ meq of $NH_3 = meq of H_2SO_4$ $76 \times 1 =$ Volume (ml) $\times N$ $\frac{76}{134} = N \implies N = 0.57 N.$ Ans Q4. Sol: CuSO₄. 5H₂O \longrightarrow Cu²⁺ ion 0.5 lit & 0.01 M : Moles of Cu^{2+} formed = Volume × Morality = $0.5 \times 0.01 = 0.005$ \therefore Moles of CuSO₄. 5H₂O = moles of Cu²⁺ = 0.005 $\frac{\text{weight CuSO}_4.5\text{H}_2\text{O}}{(63.5+96+90)} = 0.005$ wt. of CuSO₄.5H, $O = 0.005 \times (249.5) = 1.2475 g$ Ans

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

Q5. Sol: (a) $\frac{0.365g}{100} = 3.65 \,\text{gm/lit}$ weight volume $\therefore \text{ Mole/lit} = \frac{\text{weight/lit}}{\text{Mol.wt}} = \frac{3.65}{36.5} \text{M} = 0.1 \text{M}$ (b) $\frac{\text{moles}}{\text{wt}_{\text{H20}}} \frac{\frac{3}{30}}{\frac{250}{1000} \text{ kg}} = \frac{\frac{1}{10}}{\frac{1}{4}} = \frac{4}{10} = 0.4 \text{ m}.$ Ans Q6. Sol: From principle of dilution Moles of solute before = moles of solute after dilution $V_1 M_1 = (V_1 + V_2) M_2 \qquad (V_f = V_1 + V_2)$ ∴ Volume of water added $(V_2) = \frac{V_1(M_1 - M_2)}{M_2}$ Ans Q7. Sol: $Ca(OH)_2 + H_3PO_4 \longrightarrow CaHPO_4 + 2H_2U$ Since only two hydrogens are removed from H3PO4 its valancy factor is 2. \therefore Eq. wt. = $\frac{\text{Mol.wt}}{\text{V f}} = \frac{98}{2} = 49$ Ans Q8. **Sol:** $V_f = 250 \text{ ml}$ $M_{f} = 0.2 M$ $M_1 = 11.7 M, V_1 = ?$ $V_1M_1 = V_fM_f$ $V_1 \times 11.7 = 250 \times 0.2 = 250 \times \frac{2}{10} = 50$ $V_1 = \frac{50}{11.7} = 4.27 \text{ mL Ans}$ Q9. Sol: Let volume of 12 N HCl is taken x ml then volume of 3N HCl will be = (1000-x).:. From law of dilution $(1000 - x) \times 3 + x \times 12 = 1000 \times 6$ $3000 + 9x = 6000 \qquad \Rightarrow 9x = 3000$ X = 333.33 mL Volume of 12 N HCl = 333.33 mL Volume of 3N HCl = 666.67 mL Ans

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

Q11. Sol: NaOH + HNO₃ \longrightarrow NaNO₃ + H₂O 3.2 g 18.9 g 1 lit 1 lit Let V1 volume of NaOH & V2 volume of HNO3 is taken for having neutral solution meq. of NaOH = meq. of HNO₃ $V_1 \times \frac{32}{400} \times 1 = V_2 \times \frac{18.9}{63} \times 1$ $V_1 \times \frac{4}{50} = V_2 \times \frac{3}{10}$ $\frac{V_1}{V_2} = \frac{3 \times 5}{4} = \frac{15}{4} = \frac{3.75}{1}$ Ans Q12. Sol: Density = sp. gravity = 1.84 g/litV = 10 ml% by weight = 98% $\therefore \text{Normality} = \frac{\% \text{by wt} \times \text{density} \times 10}{\text{Eq.wt. H}_2 \text{SO}_4} = \frac{98 \times 1.84 \times 10}{98/2} = 36.8 \text{ N}$ $H_2SO_4 + NaOH \longrightarrow Na_2SO_4 + H_2O$ 36.8 N 2.5 N V=10 ml V=?meq. of $H_2SO_4 = meq.$ of NaOH $36.8 \times 10 = 2.5 \times V$ Volume = $\frac{36.8 \times 10}{2.5}$ = 147.2 mL Q13. Sol: Molarity = $\frac{\% \text{ by wt} \times \text{density} \times 10}{\text{Mol.wt}} = \frac{13 \times 1.09 \times 10}{98} = 1.446 \text{ M.}$ Molarity (m) = $\frac{\text{mole of H}_2\text{SO}_4}{\text{weight of solvent}}$ % by wt = $\frac{13}{100}$ wt of H₂SO₄ 13 wt of solution 100 $\Rightarrow \frac{WH_2SO_4}{WH_2O + WH_2SO_4} = \frac{13}{100}$ $W H_2SO_4 = (WH_2O + W H_2SO_4) \frac{13}{100}$

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

 $WH_2SO_4\left(1-\frac{13}{100}\right) = \frac{13}{100}WH_2O$ $\frac{WH_2SO_4}{WH_2O} = \frac{13}{87}$ $\therefore \frac{\text{wt H}_2\text{SO}_4}{98} = \frac{13}{98 \times 87}$ WH,O Let V_f is the final volume, then $V_1M_1 = V_fM_f$ $100 \times 1.446 \times 2 = Vf \times 1.5$ $Vf = \frac{144.5 \times 2}{1.5} = 193.3 \,\text{mL}$ Ans Q14. **Sol:** N_r = $\frac{1.8 \times 98 \times 10}{49} = 36.8$ N .:. From principle of dilution $V_f N_f = V_1 N_1$ $V_f = \frac{200 \times 0.5}{38.8} = 2.71 \text{ mL}$ Ans Q15. Sol: Na₂CO₃ + A \longrightarrow V=26 mL 10 mL N = 1 NFrom principle of chemical equivalence meq. of $Na_2CO_3 = meq.$ of A $26 \times 1 = 10 \times N_A$ $N_A = \frac{26}{10} = 2.6 N$ V = 26 ml 40 mm N = 1N $26 \times 1 = 40 \times N_{\rm B}$ $N_{\rm B} = \frac{26}{40} N = \frac{2.6}{4} N$ Let V₁ ml of A & V₂ ml of B is equired to make 1 lit of 1 N sol-ⁿ then $V_1 + V_2 = 1$ lit = 1000 ml-(1) Also, $V_1V_4 + V_2N_B = V_2N_2$

$$V_{1} \times 2.6 + (1000 - V_{1}) \frac{2.6}{4} = 1000 \times 1$$
$$V_{1} \times 2.6 \times \frac{3}{4} = 1000 - \frac{2.6}{4} \times 1000$$

 $\frac{\text{mole of H}_2\text{SO}_4}{\text{WH}_2\text{O(gm)}} = \frac{13}{98 \times 87}$ $\therefore \text{molaity} = \frac{\text{mole of } H_2 \text{SO}_4}{\text{wt } H_2 P(9m)} \times 1000$ $=\frac{13\times1000}{98\times87}=1.525\,\mathrm{m}$

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

 $V_1 \times \frac{2.6 \times 3}{4} = 1000 \times \frac{1.4}{4}$ $V_1 = \frac{1000 \times 1.4}{2.6 \times 3} = 179.5 \,\mathrm{mL}$ Volume of A required = 179.4 ml Volume of B required = 1000 - 179.4 = 820.6 mL Ans Q16. + $Cr_2O_7^2 \longrightarrow Cr^{3+} + Fe^{2+}$ Sol: Fe²⁺ 25 ml 32.4 ml v.f = 10.0154 M M=? (v.f = 6)meq of Fe^{2+} = meq of $Cr_2O_7^{2-}$ $25 \times M \times 1 = 32.45 \times 0.0153 \times 6$ $M = \frac{32.45 \times 0.0153 \times 6}{25} = 0.1192 M$ Ans Q17. Sol: Initial weight of HCl present in $\frac{N}{2}$ of 1 lit solution $=\frac{1}{2} \times 1 \times 1 \times 36.5 = 18.25 \,\mathrm{gm}$ weight lost = 2.675 gm : weight remained = 18.25 - 2.675 = 15.575 g volume final = 750 mL (i) $N_r = \frac{15.575 g}{36.5 g} \times 1000 = 0.569 N.$ (ii) no. of meq in 100 ml solution = $100 \times \frac{1}{2} = 50$ meq Q18. $CuSO_4 \longrightarrow ZnSO_4 + Cu$ Sol: Zn + 200 ml 10 gm Let x gm of Zn has reacted Then wt of Cu produced = $\frac{x}{65.4} \times 63.5 \text{ g}$ \therefore (mole of Zn reacted = mole of Cu produced) A/q, wt of Ξ n remained + wt of Cu produced = 9.810 g $(10-x) + \frac{63.5x}{65.4} = 9.810 \,\mathrm{g}$ $\frac{654 - 65.4x + 63.5x}{65.4} = 9.810g$ 654 - 1.9 x = 641.571.9 x = 12.426

> JEE (MAIN & ADV.), MEDICAL + BOARD By: Er. Rishi Kumar (B.Tech. IIT Kanpur) Rishi Chemistry Classes, Naya Tola , Patna -4

NEWTON CLASSES

$$x = \frac{12.426}{1.9} = 6.54$$

(i) The weight of Cu deposited = $\frac{63.5}{65.4} \times 6.54 = 6.35$ g

(ii) Molarity =
$$\frac{6.35}{\frac{63.5}{200} \times 1000} = \frac{5}{20} = 0.5 \text{ M}$$

Q19.

Q20.

Sol: (1) Cu + $Fe_2(SO_4)_3 \longrightarrow CuSO_4 + 2FeSO_4$

(If it is assumed that this is the reaction one)

0.108 g excess meq. of $FeSO_4 = meq. of CuSO_4$

$$=\frac{0.108\times1000}{63.5/2}=3.38$$

Only FeSO4 can undergo oxidation, because it is only in lower oxidation State.

(2) $FeSO_4$ + $KMnO_4 \longrightarrow Fe^{3+} + Mn^{2+}$ 33.7 0.1 N meg of $FeSO_4 = meg of KMnO_4$ $= 33.7 \times 0.1 = 3.37$ Since meq in reaction (1) = meq of reaction (2) \therefore v.f of FeSO₄ in (1) = v.f. of FeSO₄ in (2). In reaction (1) In reaction(2) $Fe^{3+} \longrightarrow Fe^{2+}$ $Fe^{2+} \longrightarrow Fe^{3+}$ So, the assumed reaction (1) is the correct one. Sol: $W(NaOH + Na_2CO_3) = 2.013 g$ Let x g of Na₂CO₃ is present in the commercial sample : wt of NaOH = (2.013 - x) g ... The amount is dissolved in 250 mL $\therefore \text{ Quartz present in 10 ml} = \left(\frac{2.031}{250}\right) \times 10 = \frac{2.031}{25} \text{g}$

: wt. of Na₂CO₃ in 10 ml =
$$\frac{x \times 10}{250} = \frac{x}{25}g$$

wt. of NaOH in 10 ml =
$$\frac{2.013 - x}{250} \times 10 = \frac{2.013 - x}{250}$$

 $NaOH + H_2SO_4 \longrightarrow Na_2SO_4 + H_2O$ (v.f.=1) $Na_2CO_3 + H_2SO_4 \longrightarrow Na_2SO_4 + H_2O + CO_2$ (v.f. = 2)meq of NaOH + meq of Na₂CO₃ = meq of H_2SO_4

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R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

 $\frac{(2.013 - x)}{\frac{25}{40/1}} \times 1000 + \frac{\frac{x}{25} \times 100}{\frac{106/2}{2}} = 20 \times 0.1$ $\Rightarrow \frac{(2.013 - x)}{25 \times 40} \times 1000 + \frac{x}{25 \times 43} \times 1000 = 2$ $\Rightarrow (2.03 - x) + 0.93 x = 2$ $\Rightarrow 0.07 \text{ x} = 2.013 - 2 = 0.013$ $x = \frac{0.013}{0.069} = 0.186$:.% of Na₂CO₃ = $\frac{x}{2.013} \times 100 = 9.26\%$ Q21. Sol: KCl + KOH + HCl \longrightarrow KCl + H₂O 120 ml 0.12 N meq. of KOH = meq of HCl(Only KOH will react with HCI). $\overline{\text{Eq.wt.of KOH}} \times 1000 = 120 \times 0.12$ $\frac{wt}{56_1} \times 1000 = 120 \times 0.12$ wt = $\frac{120 \times 0.12 \times 56}{1000} = 0.8064$ g (ii) KCl + AgNO₃ $\xrightarrow{\text{HNO}_3}$ AgCl \downarrow + AgNO₃ total $\therefore \frac{\text{wt of KCl}_{\text{total}}}{74.5/1} = \frac{3.7}{143.5}$ wt of KCl $= \frac{3.7 \times 74.5}{1415} = 1.921g$ Also wt of KCl produced from KOH = $\frac{0.8064}{56} \times 74.5 = 1.0728$ g : wt. of KCl originally present = 1.921 - 1.0728 = 0.8482 g : total wt. initially = 0.8482 + 0.8064 = 1.6546 g :.% of KOH = $\frac{\text{wt of KOH}}{\text{total Wt}} \times 100 = \frac{0.8064 \text{ g}}{1.6546} \times 100 = 48.74\%$ Ans

Q22.

Sol: 10.03 g of vinegar (contain CH₃COOH)

The solution is made upto 100 ml and out of which 25 ml is taken out for the experiment.

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

 $2CH_3COOH + Ba(OH)_2 \longrightarrow Ba(CH_3COO)_2 + 2H_2O$ V = 25 ml0.0176 M v.f = 134.30 ml v.f = 2wt of $CH_3COOH \times 1000 = 34.30 \times 0.0176 \times 2 = 1.2074$ eq.wt $\frac{\text{wt of CH}_{3}\text{COOH}}{60/1} \times 1000 = 1.2074$ wt CH₃COOH in 25 ml = $\frac{1.2074 \times 60}{1000} = 0.0724$: wt of CH₃COOH in 100 ml = $\frac{0.0724}{25} \times 100 = 0.290\%$:.% of CH₃COOH in vinegar = $\frac{0.296}{10.3} \times 100 = 2.90\%$ Ans Q23. Sol: $3Zn^{2+} + 2[4K^{+}Fe(CN)_{6}^{4+}]$ → K₂Zn₃[Fe(CN)₆]₂ → OK 1.5432 g 0.1043 M (Impure) 34.68 ml From balanced reaction coefficient (stoichiometry) 3 mole of $Zn^{2+} \equiv 2$ mole of K₄[Fe (CN)₆] $2 \times \text{mole of } Zn^{2+} = 3 \times \text{mole of } K_4 [Fe (CN)_6]$ $2 \times \frac{\text{wt}}{64.5} = 3 \times 0.1043 \times \frac{34.68}{1000}$ wt actually present = 0.35 g ∴% of Zn²⁺ = $\frac{0.35}{1.5432} \times 100 = 22.7\% = .23\%$ Q24. Sol: FeSO₄. 7H₂O + KMnO₄ \longrightarrow Fe³⁺ (v.f = 1)5.4 ml 0.1 N meq of $FeSO_4.7H_2O$ = meq. of KMnO₄ $\frac{\text{wt}}{278/1} \times 1000 = 5.4 \times 0.1$ $wt = \frac{0.54 \times 278}{1000} = 0.15g$: wt of Fe₂(SO₄)₃. 9H₂O in the sample = 5.5 - 0.15 = 5.35 g :. Moles of Fe₂(SO₄)₃. 9H₂O = $\frac{535}{562}$ = 0.0095 mole; Ans

Q25.

Sol:
$$5U^{4+} + 2MnO_4^- + 6H_2O \longrightarrow 5UO_2^{2+} + Mn^{2+} + 4H_3O^+$$

Balance reaction is given, so it is better to use Mole – stoichiometric relationship

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

5 mole of $U^{4+} = 2$ mole of MnO₄⁻ 2 mole of $U^{4+} = 5 \times \text{mole of } MnO_4^ 2 \times \frac{0.5}{238} = 5 \times \frac{50}{1000} \times M$ $M = \frac{2 \times 5 \times 100}{2380 \times 5 \times 5} = \frac{4}{238} = 0.0168 M$ Ans Q26. **Sol:** $2KH C_8H_4O_2 + Ba(OH)_2 \longrightarrow 2H_2O + 2K^+ + 2C_8H_4O_2^{2-} + Ba^{2+}$ 0.2 g 27.80 ml Mol. wt = 204.2 g2 mole of KH $C_8H_4O_2 \equiv 1$ mole of Ba(OH)₂ 1 mole of KH $C_8H_4O_2 = 2 \times \text{mole of Ba}(OH)_2$ $\frac{0.2}{204.2}$ $= 2 \times \frac{27.80}{1000} \times M$ $M = \frac{200}{204.2 \times 2 \times 27.80} = 0.0176 M \text{ Ans}$ Q27. Sol: Mg + MgO Dissolved in H₂SO₄ (125 ml & 0.1 N H2SO₄) (1) Mg + H₂SO₄ \longrightarrow MgSO₄ + H₂ \uparrow 120.1 ml At $T = 27.3^{\circ}$ C, P = 1 atm (2) MgO + $H_2SO_4 \rightarrow MgSO_4 + H_2O$ H_2SO_4 remained = 0.02 N \therefore H₂SO₄ reacted = (0.1 - 0.02) = 0.08 N. From equation(1) mole of H_2 evolved = mole of Mg = mole of H_2SO_4 used in (1) 120.1 WtM 0.821×300.3 wt of Mg = 0.117 g Mole of $H_2SO_4 = 4.871 \times 10^{-3}$ (reacted in reaction (1)) Moles of H_2SO_4 reacted in reaction (2) = Total moles reacted - moles reacted reaction (1) $=\frac{125}{1000}\times 0.08/2 - 4.871\times 10^{-3}$ $=(5-4.871) \times 10^{-3} = 0.129 \times 10^{-3}$ \therefore Moles of MgO in reaction (2) = mole of H₂SO₄ reacted in (2) $wt MgO = 0.129 \times 10^{-3}$

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

wt MgO = 0.00516 g ... Total wt (Mg - 1000) 0.1178 g + 0.00516 g = 0.123 g % by wt of Mg = $\frac{0.1170}{0.123} \times 100 = 95.9\%$ Ans Q28. **Sol:** $MH_2SO_4 \frac{1.18 \times 24.7 \times 10}{98} = 2.97 M$ Volume = 75 mLWeight of Al = 2.7 g $H_2SO_4 \longrightarrow Al_2(SO_4)_3 + H_2$ Al + 2.7-g 75 ml 2.97 M v.f=3 (v.f=2)meq of Al = meq of H_2SO_4 reacted. $\frac{2.7}{27/2} \times 1000 = \text{mmole of } \text{H}_2\text{SO}_4 \times 2$ $\frac{300}{2}$ = mole of H₂SO₄ reacted. \therefore mole of H₂SO₄ remained = 75 × 2.97 - 150 = 73 mole :. Molarity $=\frac{73}{400}=0.18$ M Ans Q29. Sol: 4 g of NaCl + Na₂CO₃ is dissolved in 250 ml 25 ml is taken for reaction Na2CO3. + \rightarrow NaCl + H₂O + CO₂ HCI V = 25 ml50 m $\frac{1}{10}$ N V. f=2 Mol. wt.= 106 meq. of $Na_2CO_3 = meq$ of HCl used $\frac{\text{wt.in 25ml}}{106} \times 1000 = 50 \times \frac{1}{10}$ 106/ wt in 25 ml = $\frac{5 \times 53}{1000}$... wt in 250 ml = $\frac{5 \times 53}{1000} \times \frac{250}{25} = 2.65 \text{ g}$:. % of Na₂CO₃ = $\frac{2.65}{4} \times 100 = 66.25\%$ % of NaCl = $\frac{1.35}{4} \times 100 = 33.75\%$ Ans

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

86

Q30. Sol: M₂CO₃ & M'₂CO₃ (suppose the two carbonates of univalent metal are) Say x gm (1 - x) gLet At. Wt. of M = 7g & M' is y gm, then meq of Metal carbonates = meq of HCl $\frac{x}{74/2} \times 1000 + \frac{1-x}{(2y+60)/2} \times 1000 = 44.4 \times 0.5 - - - - - (1)$ Also A/q, moles of $M_2CO_3 = moles of M'_2CO_3$ $\frac{x}{74} = \frac{1-x}{(2y+60)}$ $(1) \Rightarrow \frac{2x}{74} \times 1000 + \frac{2x}{74} \times 1000 = 22.2$ $\Rightarrow \frac{4x}{74} \times 1000 = 22.2 \Rightarrow x = \frac{22.2 \times 70}{4 \times 1000} = 0.41g$:. Wt of $M_2CO_3 = 0.41$ g Wt of $M_{2}CO_{3} = 0.59 g$ Also, $\frac{x}{74} = \frac{1-x}{24+60} \implies \frac{0.41}{74} = \frac{0.59}{2y+60}$ $\Rightarrow 2y + 60 = \frac{0.59 \times 74}{0.41}$ Y = 23.2 g \therefore At. Wt. of M' = 23.2 g; Ans $\frac{(1-x)}{106.4}$ × 142.0 $M_2CO_3 + M'_2CO_3 \longrightarrow M_2SO_4 + M'_2SO_4$: wt of SO₄²⁻ formed = $\frac{x}{74} \times 110 + \frac{1-x}{1064} \times 142.4$ $=\frac{0.41}{74}\times110+\frac{0.59}{1064}\times142.4$ = 0.61 g + 0.789 g = 1.4 g Ans Q31. Sol: $W(Na_2CO_3 + K_2CO_3) = 1.22 \text{ g}$ dissolved in 100 ml 20 ml extracted

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

Let wt of Na₂CO₃ is x gm, then wt of $K_2CO_3 = 1.22 - x$. In 20 ml, Wt. of Na₂CO₃ = $\frac{x}{100} \times 20 = \frac{x}{5}g$ wt of K₂CO₃ = $\frac{1.22 - x}{100} \times 20 = \frac{(1.22 - x)}{5}g$ $Na_2CO_3 + K_2CO_3 + HC1 \longrightarrow NaCl + KCl + H_2O + CO$ meq of Na_2CO_3 + meq of K_2CO_3 = meq of HCl. $\Rightarrow \frac{\overline{5}}{106/2} \times 1000 + \frac{\overline{5}}{138/2} \times 1000 = 40 \times 0.1$ $\Rightarrow \frac{x}{265} \times 1000 + \frac{(1.22 - x)}{345} \times 1000 = 4$ $\Rightarrow \frac{x}{265} + \frac{(1.22 - x)}{345} = \frac{4}{1000}$ $\Rightarrow \frac{345x - 265 - x + 265 \times 1.22}{265 \times 345} = \frac{4}{1000}$ $\Rightarrow 80 x + 323.3 = 365.7$ \Rightarrow 80 x = 42.4 $\Rightarrow x = \frac{42.4}{80} = 0.53g$: wt of Na₂CO₃ present in 1.22 g mix = 0.53 g Ans x 5×106 ×197g BaCO $Na_2CO_3 + K_2CO_3 + BaCl_2 \rightarrow$ (1.22 - x)x $\int \frac{(1.22-x)}{5\times 138} \times 197g$ 5 wt. of BaCO3 obtained $= \frac{x}{5 \times 106} \times 197 + \frac{1.22 - x}{5 \times 138} \times 197$ = (0.197 + 0.197) g = 0.397 g Ans Q32. $W(KCl + KClO_3) = 1.0g$ Sol: is dissolved in 250 mL Say (x g) (y g)25 ml extracted, which will contain $\left(\frac{x}{10}$ g KCl, $\frac{y}{10}$ g KClO₃ $\right)$ $ClO_3^- + SO_2 \longrightarrow Cl^- + SO_4^{2-}$ 25 ml v. F = 2 v.f = 6AgCl 0.1435 g

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

mole of AgCl = $\frac{0.1435}{143.5} = 0.001$ mole \therefore mole of Cl = 0.001 mole : All of the chlorine comes from ClO₃⁻ & Cl⁻ .. (mole of KCl + mole of Cl in KClO₃) in 25 ml = mole of Cl in AgCl (By POAC principle) : (mole of KCl + mole of KClO₃) in 25 ml = mole of AgCl $\frac{x}{10} + \frac{y}{10} = 0.001$ $\frac{x}{74.5} + \frac{y}{122.5} = 0.01 - - - - (1)$ Also ClO_3^- + $Fe^{2+} \longrightarrow Fe^{3+} + Cl^$ v.f = 6v.f = 1meq of ClO_3^- = meq of Fe²⁺ (used in the reac-ⁿ) $\frac{y/10}{122.5/6} \times 1000 = 30 \times 0.2 - 37.5 \times 0.08$ $\frac{y}{204.16} \times 1000 = (6-3)$ \Rightarrow Y = 0.6125 g \therefore moles of KClO₃ = $\frac{y}{122.5} = 5 \times 10^3$ moles of KCl = $\frac{x}{74.5}$ = 5×10⁻³ moles Ans Q33. Sol: MnO₂ + H₂C₂O₄ \longrightarrow Mn²⁺ + CO₂ 0.6 g 200 ml (impure) 10 N Let x gm of MnO2 of present in the reaction, then meq of $MnO_2 = meq$ of $H_2C_2O_4$ (reacted) $\frac{x}{87/2}$ ×1000 = meq of H₂C₂O₄(reacted) meq of H₂C₂O₄ (remained) in 500 ml = $200 \times \frac{1}{10} - \frac{2x \times 1000}{87} = 20 - \frac{2000x}{87}$ $H_2C_2O_4 + KMnO_4 \longrightarrow Mn^{2+} + CO_2$ remained 50 ml 1/30 N in 100 ml

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

meq of $H_2C_2O_4$ in 100 ml = meq of KMnO₄ used. $\frac{20 - \frac{2000 \text{ x}}{87}}{5} = 50 \times \frac{1}{30} = \frac{5}{3}$ $\frac{2000x}{87} = 20 - \frac{25}{3} = \frac{60 - 25}{3} = \frac{35}{3}$ $x = \frac{35 \times 87}{3 \times 2000} = 0.5075 \,\mathrm{gm}$:. % of MnO₂ in the sample of ore = $\frac{x}{0.6} \times 100 = \frac{0.5075}{0.6} \times 100 = 84.58\%$ Ans Q34. Sol: In pressure of bromocresol green $H_3PO_4 + nOH \longrightarrow nH_2O + [H_{(3-x)}PO_4]^{n-1}$ 23.1 ml 25 ml 0.107 M 0.115 M V.f = 1V. f = n $\{3-(3-n)\} = no. of replaced hydrogen$ meq of $H_3PO_4 = meq$ of NaOH $25 \times 0.107 \times n = 23.1 \times 0.115 \times 1$ $n = \frac{23.1 \times 115}{25 \times 0.107} = 1$ Ans In pressure of phenolphthalein indicator $H_3PO_4 + nOH^- \longrightarrow n'H_2O + [H_{(3-n')}PO_4]^n$ 46.8 ml 25 ml 0.115 M 0.107 M v. f = n'v.f = 1meq of $H_3PO_4 = meq of OH$ $25 \times 0.107 \times n' = 46.8 \times 0.115$ $n^2 = \frac{46.8 \times 0.115}{25 \times 0.107} = 2$ Ans Q35. Sol: FeSO4(NH4)2 SO4. xH2O 9.8 g in 250 ml 20 ml extracted wt. of FeSO₄(NH₄)₂SO₄.xH₂O present in 20 ml of solution = $\frac{9.8}{250} \times 20 = 0.784$ g $FeSO_4 (NH_4)_2SO_4.xH_2O + KMnO_4 \longrightarrow Mn^{2+} + Fe^{3+}$ 3.53 g/ lit 0.784 g 90% pure v f = 120 ml

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meq of FeSO₄(NH₄)₂SO₄. xH₂O = meq of KMnO₄ $\frac{0.784}{\left(\frac{284+18x}{1}\right)} \times 1000 = \left(\frac{3.53 \times \frac{90}{100}}{158/5}\right) \times 1000 \times \frac{20}{100}$ $\frac{0.784}{(284+18x)} = \frac{0.1}{50}$ 284 + 18x = $\frac{0.784 \times 50}{0.1} = 7.84 \times 50 = 392$ 18 x = 392 - 284 x = $\frac{108}{18} = 6$ Ans

Q36.

Sol: 10 ml of Solution containing 10.25 g of H₂O₂ diluted to 250 ml, which will contain same mass of H₂O₂

Now 25 ml is extracted which will contain $=\frac{10.25}{250} \times 25 = 1.025 \text{ g}$ H_2O_2 + . $KMnO_4 \longrightarrow O_2 + Mn^{2+}$ 1.025 g 35.8 ml (v.f=2) $\frac{1}{10}N$ meq of H_2O_2 in 25 ml = meq of KMnO₄ used. $\frac{\text{wt}}{34/2} \times 1000 = 35.8 \times \frac{1}{10} = 3.58$ wt of H₂O₂ in 25 ml = $\frac{3.58 \times 17}{1000} = 0.06086$ g :. wt of 250 ml (Diluted) = $0.06086 \times 10 = 0.6086$ g ...wt of original 10 ml solution = 0.6086 g \therefore wt in original 100 ml solution = 0.6086 × 10 g = 6.086 g Volume strength = Normality of solution × 5.6 $= \frac{6.086/17}{100} \times 1000 \times 5.6 \ [:.5.6 \text{ N H}_2\text{O}_2 \text{ solution} = 1 \text{ volume of H}_2\text{O}_2 \text{ solution}]$ = 20 volume Ans Q37. Sol: $H_2O_2 + KI \longrightarrow H_2O + I_2$ 50 ml (excess) (v.f. = 2)meq of I_2 librated = meq of H_2O_2 -----(1) $I_2 + Na_2S_2O_3 \longrightarrow Na_2S_4O_6 + 2NaI$ (v.f = 2)

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meq of $I_2 = meq of Na_2S_2O_3$ -----(2) Since v.f. of I₂ in both (1) & (2) are same, so (1) & (2) \Rightarrow meq of $H_2O_2 = meq$ of $Na_2S_2O_3$ $50 \times N = 20 \times 0.1$ $N = \frac{2}{50} = \frac{4}{100} = 0.04 N$ Also, $\frac{\text{gm.eq}}{\text{lit}} = N = 0.04 \text{ N}$ $\frac{\text{wt}/17}{\text{lit}} = 0.04 \text{ N} \implies \frac{\text{wt}}{\text{lit}} = 0.04 \times 17 = 0.68 \text{ g/lit} \text{ Ans}$ **O38**. Sol: 3.546 g dissolved in bleaching powder [Co(OCl)Cl] $Ca(OCI)CI + KI \longrightarrow I_2 + Cl_2$ v.f. = 225 ml (v.f. = 2)meq, of Ca(OCl)Cl = meq of I_2 evolved -----(1) Also, $I_2 + Na_2S_2O_3 \longrightarrow Na_2S_4O_6 + 2NaI$ (v.f. = 2)meq of I_2 (reacted) = meq of $Na_2S_2O_3$ -----(2) $(1) \& (2) \Rightarrow .$ Meq of Ca(OCl)Cl = meq. of Na₂ $\$_2O_3$ $= 20 \times 0.125$ $\frac{\text{wt of Ca(OCI)Cl}}{127/2} \times 1000 = 20 \times 0.125$ wt of Ca (OCl)Cl present in 25 ml = 0.15875 g wt of Cl_2 evolved $\times 1000 = 20 \times 0.125$ (meq of $Cl_2 = meq$ of $I_2 = meq$ of hypo) wt of Cl₂ evolved from 25 ml = $\frac{20 \times 0.125 \times 35.5}{1000} = 0.08875$ g \therefore wt of Cl₂ evolved from 100 ml sol-ⁿ = $\frac{0.8875}{25} \times 100 = 0.355$ $\therefore \% \text{ available chlorine} = \frac{0.355}{3.546} \times 100 = 10.01\%$ Ans Q39. $CuSO_4 + Zn \longrightarrow ZnSO_4 + Cu \downarrow$ Sol: 25 g 100 ml Initially: 1 M After reaction 0.8 M \therefore Morality of SO₄²⁻ after reaction = 0.8 + 0.2 = 1.0 M

OFFICE AT $\boldsymbol{\varsigma}$ 606,6th FLOOR RIOM TOWFR NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD (SO₄²⁻ ion comes from ZnSO₄ & CuSO₄ both) Also, moles of Zn consumed = moles of CuSO₄ reacted wt.consumed 100×0.2 65.4g 1000 wt. consumed = $6.54 \text{ g} \times 0.2 = 1.308 \text{ g}$ \therefore wt. of Zn rod after reaction = (25 - 1.308) g = 23.692 g Ans (b) $Cu + ZnSO_4 \longrightarrow No$ reaction (Cu is very less electropositive than Zn so can't replace Zn from $ZnSO_4$) 25 g 2M Zn^{2+} after reaction = 2M Ans Q40. Sol: In the case of phenolphthalein as indicator only Na2CO3 reacts with HCl to give NaHCO3 $Na_2CO_3 + HCl$ \longrightarrow NaHCO₃ + NaCl 25 ml 12 ml & $\frac{1}{20}$ N mmole of Na₂CO₃ in 25 ml = mmole of HCl = $12 \times \frac{1}{20} = 0.6$ $\frac{\text{wt}}{106} \times 1000 = 0.6$ Wt Na₂CO₃ in 25 ml = $\frac{0.6 \times 106}{1000}$ = 0.0636 g :.wt Na₂CO₃ in 1 lit = $\frac{0.0636}{25} \times 1000 \neq 2.544$ g Ans In presence of methyl orange as indicator $Na_{2}CO_{3} + 2HCI \longrightarrow NaCl + H_{2}O + CO_{2}$ $NaHCO_{3} + HCI \longrightarrow NaCl + H_{2}O + CO_{2}$ mmole of Na_2CO_3 + meg of $NaHCO_3$ = meg of HCl $2 \times \text{mmole of Na}_2CQ_3 + \text{mmole of NaHCO}_3 = \text{mmole of HCl}$ $2 \times 0.6 + \text{mmole of NaHCO}_3 = 30 \times \frac{1}{20} \times 1$ mmole of NaHCO₃ = 1.5 - 1.2 = 0.3 $\therefore \frac{\text{wt of NaHCl}_3}{84} \times 1000 = 0.3$ wt of NaHCO₃ in 25 ml = $0.3 \times \frac{84}{1000} = 0.0252$:. wt. of NaHCO₃ in 1 lit = $\frac{0.0253}{25} \times 1000 = 1.089$ g Ans Q41. Sol: In presence of phenolphthalein as indicator (1) $Na_2CO_3 + HCl \longrightarrow NaHCO_3 + NaCl$

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(2) NaOH + HCl \longrightarrow NaCl + H₂O

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

meg of Na₂CO₃ + meg of NaOH = meg of HCl used mmole of Na₂CO₃ + mmole of NaOH = $17.5 \times \frac{1}{20}$ -----(1) (Because v.f. of Na₂CO₃ & NaOH = 1, in the above reaction). (1) \Rightarrow Also mmole of NaHCO₃ formed = mmole of Na₂CO₃ -----(2) Also in presence of Methyl orange NaHCO₃ $\xrightarrow{1 \text{HCl}}$ NaCl + H₂O + CO₂ 2.5 ml N/10 ∴ meq of NaHCO3 = meq of HCl mmole of NaHCO₃ = $2.5 \times \frac{1}{10} = 0.25$ \therefore (2) \Rightarrow mmole of Na₂CO₃ = 0.25 $\therefore \frac{\text{wt}}{106} \times 1000 = 0.25$ Wt. of Na₂CO₃ = $\frac{0.25 \times 106}{1000}$ = 0.0265 g Ans (1) \Rightarrow mmole of NaOH = 1/75 = 0.25 = 1.5 $\frac{\text{wt NaOH}}{40} = \frac{1.5 \times 4}{100} = 0.06 \text{ g}$ Ans Q42. Sol:- Let the chloride is MClm 0.2 g 0.47 g $\text{meq of MCl}_{\text{m}} = \text{meq of AgCl}$ $\frac{0.29}{\text{Eq.wt. MCl}_{\text{m}}} \times 1000 = \frac{0.47 \text{ g}}{\text{Eq.wt. AgCl}} \times 1000$ Eq. wt MClm = $\frac{0.2 \times 143.5}{0.47} = 61.06 \text{ g}$ Eq. wt of M + Eq. wt of Cl = 61.06 g Eq. wt of M = 61.06 - 35.5 = 25.56 g Ans Q43. Sol: (1) 4 Fe^{3+} + N₂H₄SO₄ \longrightarrow N₂ + 4F e^{2+} + 4H⁺ (v.f = 4)(2) $MnO_4^{-} + 5Fe^{2+} \longrightarrow Mn^{2+} + 5Fe^{3+} + 4H_2O$ In 10 ml solution :-From reaction (1) meg of Fe^{3+} = meg of N₂H₄SO₄ = Fe^{2+} formed -----(1) From reaction (2) meq of Fe^{2+} = meq of MnO₄⁻ (used) -----(2)

(1) & (2) \Rightarrow meq of N₂H₄SO₄ = meq of MnO₄ (because V.F Fe²⁺) $=\frac{\text{wt}}{128/4} \times 1000 = 20 \times \frac{1}{50} \times 5 = 2$ wt of N₂H₄SO₄ in 10 ml = $\frac{2 \times 32}{1000} = 0.064$ g $N_2H_4SO_4$ in 100 ml = 0.64 g \therefore wt N₂H₄SO₄ in 1 lit = $\frac{0.64 \times 1000}{100}$ = 6.4g Ans Q44. Sol: 5 g present in 500 ml of which 20 ml extracted $Ca(OCI)CI + KI \xrightarrow{CH_3COOH} I_2 + Cl_2$ 20 ml (v, f=2)meq of $Cl_2 = meq$ of $I_2 - (1)$ Also meg of $I_2 = meg$ of hypo solution used -(2) $=20 \times \frac{1}{10} = 2$ $\therefore \text{ meq of } Cl_2 = 2 \implies \frac{\text{wtCl}_2}{71/2} \times 1000 = 2 \implies \text{Wt } Cl_2 = \frac{71}{1000} = 0.071 \text{g}$ In 20 ml wt of Ca(OCl)Cl present = $\frac{5}{500} \times 20 = 0.2$ g :. % available chlorine = $\frac{0.671}{0.2} \times 100 = \frac{7.1}{0.2} = 35.5\%$ Ans Q45. (1) $Cu_2S + CuS + MnO_4 \rightarrow Cu^{2+} + SO_2 + Mn^{2+}$ Sol: (v.f = 8)200 ml & 0.75 M wv.f=5(2) MnQ₄ (remained) + Fe²⁺ \longrightarrow Fe³⁺ + Mn²⁺ (V.f=5) 175 ml & 1 M v f = 1meq of $Cu_2S \neq meq$ of CuS = meq of MnO_4 (used) mmole of $Cu_2S \times 8 =$ mmole of $CuS \times 6 = (200 \times 0.75 \times 5 - 175 \times 1 \times 1)$ Let x g of Cus is present, then wt of $Cu_2S(10 - x)g$ $\frac{7640 + 190x}{159 \times 95.5} = 0.575$ $\therefore \frac{10 - x}{159} \times 1000 \times 8 + \frac{x}{95.5} \times 1000 \times 6 = 575$ $\frac{8(10-x)}{159} + \frac{6x}{95.5} = 0.575$ 190 x = 8731.08-7640 = 1-91.0875X = 5.74 g $\frac{7640 - 7.64x + 954x}{159 \times 95.5} = 0.575$:.% of CuS = $\frac{x}{10} \times 100 = 57.4\%$ Ans

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046. Sol: 0.518 g sample meg of $MnO_4^- = meg of CaC_2O_4$ $400 \times 0.25 = 2 \times \text{mmole of } CaC_2O_4$ Mm mole of CaC₂O₄ = $\frac{40 \times 25}{2 \times 100} = \frac{10}{2} = 5$ \therefore mmole of Ca = mmole of CaO = 5 (By POAC) $\therefore \frac{\text{wt of CaO}}{56} \times 1000 = 5$ wt CaO = $\frac{56 \times 5}{1000}$ = 0.28g \therefore % of CaO = $\frac{0.28}{0.518} \times 100 = 54.1\%$ Ans Q47. Sol: (1) $\overset{+5}{B}rO_{3}^{-} + \overset{+4}{Se}O_{3}^{2-} + H^{+} \longrightarrow \overset{+6}{Se}O_{4}^{2-} + Br_{2}$ $20 \text{ ml} \quad \frac{1}{60} \text{ M}$ (v.f=5) (v.f=2) (2) $\overset{+5}{BrO_3} - \overset{+3}{A} \overset{+3}{SO_2} - + H_2O \longrightarrow \overset{+}{Br} + \overset{+5}{A} \overset{+5}{SO_4} - H_2O \longrightarrow \overset{+}{Br} + \overset{+5}{A} \overset{+5}{SO_4} + H_2O \longrightarrow \overset{+}{Br} + \overset{+5}{Br} + H_2O \longrightarrow \overset{+}{Br} + \overset{+5}{Br} + H_2O \longrightarrow \overset{+}{Br} + H_2O \overset$ (excess) 5.1 ml $(v.f=6) = \frac{1}{25}M$ (v.f = 2)(1) \Rightarrow meq of BrO₃ (reacted) = meq of SeO₃² $5 \times \text{mmole of BrO}_3$ (used) = 2 × mmole of SeO₃²⁻ -----(1) $(2) \Rightarrow \text{meq of BrO}_3$ (excess) = meq of AsO₂ used. $6 \times m$ mole of BrO₃ (excess) = 2 × mmole of ASO₂ used -----(2) Also mmole of BrO₃ (excess) + mole of BrO₃ (used in (1) = $20 \times \frac{1}{60} = \frac{1}{2}$ $(\frac{1}{2} \text{ m mole of BrO}_3 \text{ excess}) = 2 \times \text{mmole of seO}_3^{2-1}$ $\Rightarrow 5 \times \left(\frac{1}{3} - \frac{2 \times 5.1 \times \frac{1}{25}}{6}\right) = 2 \times \text{m mole of SeO}_3^{2-}$ $\Rightarrow \text{ mmole of BrO}_3 \text{ excess} = \frac{2 \times 5.1 \times \frac{1}{25}}{6} \Rightarrow 5\left(\frac{1}{3} - 0.068\right) = 2 \times \frac{\text{wt}}{(79 + 16 \times 3)} \times \frac{1}{(79 + 16 \times 3)}$ $\Rightarrow 0.6633 = \frac{\text{wt}}{127} \times 1000 \Rightarrow \text{Wt} \text{SeO}_3^{2-} = \frac{127 \times 0.6633}{1000} = 0.084 \text{g}$ Note:- Here v.f. of BrO3 in both reaction are different, so meq. of BrO3 can't equated.

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Q48.

Sol: Normality of concentrated HCl solution = $\frac{37 \times 1.19 \times 10}{36.5}$ = 12.06 N

By principle of dilution $V_1N_1 = V_fN_f$

 $4.5 \times 12.06 = 100 \times N_f$ & $N_f = \frac{4.5 \times 12.6}{100} = 0.543 N_f$

HCl + AgNO₃ \longrightarrow AgCl + HNO₃ 10 ml 0.108 M 0.543 N V = ? For complete precipitation meq of HCl = meq of AgNO₃ 10 \times 0.543 = 0.108 \times VAgNO₃ VAgNO₃= 50.26 ml Ans

Q49.

Sol: NaKC₄H₄O₆.4H₂O $\xrightarrow{\Delta}$ NaKCO₃ By POAC mmole of NaKCO₃(produced) = mmole of NaKC₄H₄O₆. 4H₂O -(1) NaKCO₃ + H₂SO₄ \longrightarrow NaKSO₄ + H₂O + CO₂ 41.72 0.1307 N excess H₂SO₄ required = 1.91 ml of 0.1297 N NaOH

meq of NaKCO₃ = meq of H₂SO₄ used = $(41.72 \times 0.1307 - 1.91 \times 0.1297)$ mmole of NaKCO₃ × 2 = 5.21

(1) \Rightarrow mmole of NaKC₄H₄O₆ 4H₂O = $\frac{1}{2}$

$$\Rightarrow \frac{\text{wt}}{282} \times 1000 = \frac{5...}{2} \Rightarrow \text{Wt} = 0.73361$$

$$\therefore \% \text{purity} = \frac{0.73461}{0.9546} \times 100\% = 76.9\% \text{ Ans}$$

Q50.

Sol: KMnO₄ + K₂Cr₂O₇ + KI $\xrightarrow{H^+}$ I₂ + Cr³⁺ + Mn²⁺ v.f = 5 v.f = 6 (v.f = 1) v.f = 2 Also I₂ + hypo \longrightarrow Γ + S₄O₆²⁻ (v.f = 2) (Na₂S₂O₃) \therefore meq of KMnO₄ + meq of K₂Cr₂O₇ = meq of I₂ = meq of hypo solution = 60 × 0.1 = 6 mmole of KMnO₄ × 5 + mmole of K₂Cr₂O₇ × 6 = 6 $\frac{\text{wt KMno_4}}{158} \times 5 \times 1000 + \frac{(0.24 - \text{wt}_{\text{KMno_4}})}{294} \times 6 \times 1000 = 6$

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wt $KMnO_4 = x$ (say) wt $K_2Cr_2O_7 = (0.24-x) y$ 5x (0.24 - x) = 6158 294 1000 1470x - 948x + 227.52 =6 158×294 1000 552x = 279.712 - 227.52 = 52.19 $x \frac{51.19}{552} = 00.937$ wt of KMnO₄ = 0.0937 g wt of $K_2Cr_2O_7 = 0.146$ g wt of Cr = $\frac{0.146}{294} \times 2 \times 52 = 0.051$ & wt of Mn = $\frac{0.0937}{158} \times 55 = 0.034$ % of Cr = $\frac{0.051}{0.24} \times 100 = 20.8\%$ & % of Mn = $\frac{0.034}{0.24} \times 100 = 14.17\%$ Q51. Sol: 50 ppm = 50 ml NH₃ per $[O^6 ml of air$ Density NH₃ at Room temp = 0.771 g / lit $NH_3 + HCl \longrightarrow NH_4Cl$ 100 ml 0.0105 M In 10 min, volume of air passed = 10 lit / min × 10 min = 100 lit air \therefore meg of NH₃ present in 100 lit air = meg of HCl reacted = $100 \times 0.105 - 13.1 \times 0.058$ mmole of NH_3 in 100 lit air = 0.2797 $\frac{\text{wr}}{17}$ in 10⁵ ml air = 0.2797 (a) wt of NH₃ drawn to acid solution = $0.2797 \times 17 = 4.755$ g Ans wt of NH₃ in 10^5 ml air = 0.2797 × 10 × 17 = 47.55 g :. Volume of NH₃ present = $\frac{\text{wt}}{\text{density}} = \frac{47.55}{0.771} = 61.6 \text{ ml}$ (b) ... ppm of NH₃ = 61.6 ppm Ans (c) No, because max-^m allowed ppm is 50 ppm

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Objective Questions Q1: Ans-(a) Normality = $\frac{\text{no.of equivalent}}{\text{no.of equivalent}}$ litre of solution Q2: Ans -(a) $2\overset{2}{S}_{2}O_{3}^{2-}+I_{2} \longrightarrow \overset{2.5}{S}_{4}O_{6}^{2-}+2I^{-}$ $V \cdot f = 0.5 \times 2 = 1$ \therefore eq. Wt. = $\frac{Mol.wt}{wf}$ = Mol.wt Q3: Ans -(c) $\overset{+2}{V}$ + Fe₂O₃ \longrightarrow FeO + $\overset{+5}{V}_2$ O₅ $v.f = 3 \times 2 = 6$ \therefore eq. wt = $\frac{\text{Mol.wt}}{6}$ Q4: Ans -(b) $Na_2CO_3 + HCl \longrightarrow NaHCO_3 + NaCl$ v.f = 1, because it reacts with one mole of H $\therefore \text{ eq.wt} = \frac{\text{Mol.wt}}{\text{wf}} = \frac{106}{1} = 106$ Q5: Ans -(b) v.f. of I_2 in $I_2 + 2S_2O_3^{2-} \longrightarrow S_4O_6^{2-} + 2I$ is 2 \therefore eq. wt = $\frac{Mol.wt}{2}$ Q6: Ans -(c) $\operatorname{CrO_4^{2-}} \xrightarrow{H^+} \operatorname{Cr}^{3+}$ (in acidic med-") V f = 3 $\therefore \text{Eq. Wt.} = \frac{\text{Mol.wt}}{3}$ Q7: Ans -(b) eq.wt = $\frac{\text{Mol.wt}}{\text{v.f}} = \frac{158}{1}$ v.f in $MnO_4^- \longrightarrow MnO_4^{2^-}$ V.f = 1Q8: Ans -(d) meq of acid = meq of base $\frac{126}{M.W} = 20 \times 0.1 \implies Mol.wt = \frac{126}{2} = 63$ Q9: Ans -(b) NaH2PO4 . \longrightarrow Na₃PO₄ v.f = 2 because 2 H⁺ are removed from NaH₂PO₄ : meq of acid = meg of base $\frac{12}{120/2} \times 1000 = V \times 1 \times 1$ $\frac{100}{5} = V \implies V = 20$ ml Q10: Ans -(b) meq of acid = meq of base $\frac{2}{40} = \frac{3}{\text{Eq.wt}}$

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

Eq. wt of acid 60 g Q11: Ans -(b) 1 mole of HNO3 when take 4 mole of e-, oxidation state of nitrogen decrease by 4. In HNO₃, oxidation state of N = +5Final oxidation state of N = +1. Which is in N₂O. (Also it should be $\frac{1}{2}$ mole be present after reaction). Q12: Ans -(c) $N = \frac{\% \text{ by wt} \times \text{density} \times 10}{\text{Eq. wt}} = \frac{1 \times 10 \times 1}{\frac{98}{2}}$ (... For very dilute solution we can take) d solution \approx d solvent = d_w = 1 g/cc Q13: Ans - (a) $0.1 \times V = \frac{6.3}{6.3} \times 1000$ V = 1000 ml = 1 litQ14: Ans -(d) Let V is the volume required to make it deci normal Q15: Ans -(b) meg of $H_2O_2 = meg of KMnO_4$ $\frac{W}{34/2}$ ×1000 = 10×1 (because v.f of H₂O₂ = 2) $w = \frac{170}{1000} = 0.17 g$...% by wt = 85% Q16: Ans -(c) meg of NaOH = $100 \times 0.5 = 50$ meg of $H_2SO_4 = 10 \times 3 = 30$, meq of HCl = $20 \times 1 = 20$.: meg of base (NaOH) = meg of acid = meg of H2SO4 + meg of HCl : the sol-" will be neutral. Q17: Ans -(c) Normality = Molarity ×v.f (a) V . $f H_2SQ_4 = 2$ (b) V . $f H_3PO_3 = 2$ Normality will be higher if v.f will be larger (c) V.f. $H_3PO_4 = 3$ (d) V.f. $HNO_3 = 1$ as acid Q18: Ans -(d) meq. of metal = gm eq. of acid eq.wt (metal) = x g Q19: Ans -(b) $M = \frac{98 \times 1.8 \times 10}{98} = 18$ Q20:Ans -(c) meq of Na₂CO₃. X H₂O in 20 ml = meq of HCl (used) $\left(\frac{0.7}{106+18x}\right) \times \frac{20}{100} = 19.8 \times 0.1$ Calculating the value of x, we have X = 2Q21: Ans -(b) 1 N H₂SO₂ sol-ⁿ 5.6 volume of H₂O sol-ⁿ

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

100

: 5.6 volume of H₂O₂ has normality 1 N $\frac{1}{56}$ $\frac{20}{5.6} = 3.58$ Q22: Ans -(d) $V_1 N_1 = V_f N_f$ $8.3 \times 36 = (441.7 + 8.8) \times Nf$ $Nf = \frac{8.3 \times 36}{999.0} = 0.29 N = 0.3 N$ Q23: Ans -(a) meg of HCl = meg of ABCl precipitated $10 \times N = \frac{0.1435 \times 1000}{143.5/1} = 1$ $N = \frac{1}{10} = 0.1$ Q24: Ans -(b) NO3 will remain in the solution which has now volume =1 lift $\therefore \frac{500}{1000} \times 0.1 = 1 \times \text{Nf} \implies \text{Vf} = \frac{0.1}{2} = 0.05 \text{ N}$ **Q25:** Ans -(a) Ratio of amount = $\frac{\text{Amount of H}_2\text{S for ppt. of AgNO}_3}{\text{Amount of H}_2\text{S for ppt. of CuSO}_4}$ For same, moles of AgNO3 & CuSO4 present in the solution; Meq of H_2S required = meq of AgNO₃ = mmole of $AgNO_3 \times 1$ meq of H_2S required = meq of $CuSO_4$. = mmole of $CuSO_4 \times 2$ $\therefore \text{ Ratio of amount of } H_2S = \frac{\text{mmole of } AgNO_3 \times 1}{\text{mmole of } CuSO_4 \times 2} = \frac{1}{2}$ Q26: Ans -(d) 2Na₃PO₄ + 3BaCl₂ \rightarrow Ba₃(PO₄)₂ + 6NaCl 0.5 mole 0.2 mole 2 mole of Na₃PO₄ reacts with 3 mole of BaCl₂ 0.3 mole of BaCl₂ :.0.5 mole of BaCl2 is given in excess amount Na₃PO₄ is the limiting reagent \therefore mole of Ba₃(PO₄)₂ formed = $\frac{0.2}{2}$ = 0.1 mole Q27: Ans -(b) meg of acid = meg of base (caustic potash (KOH) $\frac{0.45 \times 1000}{20/v_{\rm f}} = 20 \times 0.5$ $\frac{450}{90} \times v.f = 20 \times \frac{5}{10}$ v.f =2, basicity of acid =2 Q28: Ans -(d) $1 \times 18 = 100 \times M$

R. K. MALIK' S NEWTON CLASSES JEE (MAIN & ADV.), MEDICAL + BOARD

M = 0.18 M∴ N = 0.18 × v.f = 0.18 × 2 = 0.36 N Q29: Ans -(d) For weak acid & strong base titration, final solution will be basic i.e, PH > 7 ∴ only those indicators can be used which shown and point at PH > 7, So (d) is the correct option Q30: Ans -(b) Eq of A = Eq. of B= Eq. of AxBy Q31: Ans -(b) Volume strength = 1.5 × 5.6 = 8.4 volume Q32: Ans -(a):- mole × v.f. = mole of FeSO₄ × V.f. mole of KMnO₄ × 5 = 1 × (1+2) mole of KMnO₄ × 5 = 1 mole of SO₃²⁻ × 2 (SO₃²⁻ → SO₄²⁻) ∴ mole of KMnO₄ = $\frac{2}{5}$

Q 34: Ans -(b) In disproportionation reaction, same substance oxidises as well as reduces. In (b) Cl₂ is reducing to HCl and oxidising to HClO