

**MAINS+ADVANCED
TOPIC**
**IONIC
EQUILIBRIUM**
SOLUTIONS
IONIC EQUILIBRIUM
Exercise-01

9. $\text{pH} = 1$ $\text{pH} = 2$
 $[\text{H}^+] = 0.1$ $[\text{H}^+] = 0.01$
 $V = 50$ $V = 50$
- $[\text{H}^+]$ of mixture is $[\text{H}^+] = \frac{N_1 V_1 + N_2 V_2}{V_1 + V_2} \Rightarrow \frac{50(0.1 + 0.01)}{100}$
- $[\text{H}^+] \Rightarrow \frac{0.11}{2} \Rightarrow 0.055$
 $\text{pH} = 1.26$
11. $\text{pH} = 7$ $[\text{H}^+] = 10^{-7}$, $[\text{OH}^-] = 10^{-7}$
new pH after addition of base
 $\text{pH} = 12$ $[\text{H}^+] = 10^{-12}$
 $[\text{OH}^-] = 10^{-2}$
 $[\text{OH}^+]$ concentration increase 10^5 times.
13. Relative strength = $\sqrt{\frac{K_{a_1}}{K_{a_2}}} = \sqrt{\frac{3 \times 10^{-4}}{1.8 \times 10^{-5}}} \Rightarrow 4 : 1$
16. $\text{HCOOH} + \text{KOH}$
 $40\text{mL}, 0.5(\text{M}) \xrightarrow{50\text{mL}, 0.2\text{M}} \text{HCOOH} + \text{H}_2\text{O}$
after reaction is forms Buffer solution
- $[\text{HCOOH}] = \frac{10}{90}$ $[\text{HCOOK}] = \frac{10}{90}$
- $\text{pH} = \text{p}K_a + \log \frac{[\text{salt}]}{[\text{acid}]}$ $\text{pH} = \text{p}K_a$
- $\text{pH} = 4 - \log (1.8)$ $\text{pH} = 3.75$
19. Let weak acid is HA its sodium salt is NaA
- $K_a = \frac{K_w}{K_H}$ $K_H = CH^2 \Rightarrow 0.1 \times (0.03)^2$
- $K_a = \frac{10^{-14}}{9 + 10^{-5}}$ $K_a \approx 1 \times 10^{-10}$
22. $\text{CH}_3\text{COOH} + \text{NaOH} \rightarrow \text{CH}_3\text{COONa} + \text{H}_2\text{O}$
 $100 \text{ mL}, 0.4 \text{ M}$ $100 \text{ mL}, 0.2\text{M}$
 40 20
- After reaction
- $[\text{CH}_3\text{COOH}] = \frac{20}{250}$, $[\text{CH}_3\text{COONa}] = \frac{20}{200}$
- $\text{pH} = \text{p}K_a + \log \frac{[\text{salt}]}{[\text{acid}]}$

24. $\text{pH} = \text{p}K_a$ $[\text{H}^+] = K_a = 1.8 \times 10^{-5}$
 $\text{CH}_3\text{COONa} + \text{HCl} \rightarrow \text{CH}_3\text{COOH} + \text{NaCl}$
at equivalence the $[\text{CH}_3\text{COOH}] = \frac{20}{200} \Rightarrow 0.1$
- $\text{pH} = \frac{1}{2}\text{p}K_a - \frac{1}{2}\log C$
 $\text{pH} = \frac{1}{2}[5 \log 2 - \log 10^{-5}]$
 $\text{pH} = \frac{1}{2}[6 - \log 2] \Rightarrow \text{pH} = 3 - \log \sqrt{2}$
28. $\text{pH} = \text{p}K_a + \log \frac{\text{HCO}_3^-}{\text{H}_2\text{CO}_3}$
- $7 = 7 - \log + \log \frac{\text{HCO}_3^-}{\text{H}_2\text{CO}_3} \frac{\text{HCO}_3^-}{(\text{H}_2\text{CO}_3)} = 4$
- % $\text{HCO}_3^- = \frac{4}{5} \times 100 \Rightarrow 80 \%$
30. $2\text{NaOH} + \text{H}_3\text{PO}_4 \rightarrow \text{Na}_3\text{PO}_3$
 $50\text{mL}, 0.1\text{M}$ $60\text{mL}, 0.15$
 0 7.33
- $\text{H}_3\text{PO}_4 \rightleftharpoons \text{H}^+ \text{H}_2\text{PO}_4^-$
[mass H^+ obtain from first ionization of H_3PO_4]
- $\text{pH} = \frac{1}{2}\text{p}K_a - \frac{1}{2}\log C = \frac{1}{2}\text{p}K_a - \frac{1}{2}\log C$
 $= 1.5 + 1.17 = 2.67$
32. $\text{Mg}(\text{NO}_3)_2 + 2\text{NaF} \rightarrow \text{MgF}_2 + 2\text{NaNO}_3$
 2.5 5
 0 0 2.5
- $(\text{MgF}_2) = \frac{2.5}{35}$
- $\text{MgF}_2 \rightarrow \text{Mg}^{+2} + 2 \text{ F}^-$
Ionic product
41. $\text{AgCNS} \rightleftharpoons \text{Ag}^+ + \text{CNS}^-$
 $(1-\delta_1)$ $(\delta_1 + \delta_2) \quad \delta_1$
 $\text{AgCl} \rightleftharpoons \text{Ag}^+ + \text{Cl}^-$
 $(1-\delta_2)$ $(\delta_1 + \delta_2) \quad \delta_2$
 $(K_{sp})_{\text{AgCNS}} = (\delta_1)(\delta_1 + \delta_2) \quad \dots \dots \dots \text{(i)}$
 $(K_{sp})_{\text{AgCl}} = (\delta_2)(\delta_1 + \delta_2) \quad \dots \dots \dots \text{(ii)}$

add equation (i) and (ii)

$$(\delta_1 + \delta_2)^2 = (K_{sp})_{AgCNS} + (K_{sp})_{AgCl}$$

$$(\delta_1 + \delta_2) = \sqrt{(K_{sp})_{AgCNS} + (K_{sp})_{AgCl}} = \sqrt{1.0 \times 10^{-12} + 1.7 \times 10^{-10}}$$

$$(AgT) = (\delta_1 + \delta_2) = 1.3 \times 10^{-5} \text{ divide eq. ii from i}$$

$$\frac{[Cl^-]}{[CNS^-]} = \frac{\delta_2}{\delta_1} = \frac{1.7 \times 10^{-10}}{1.0 \times 10^{-12}} \Rightarrow 1.7 \times 10^2$$

47. $IP > K_{sp}$ I.P. = $(Ca^{+2})(F)^2$
 $I.P. \Rightarrow (10^{-2})(10^{-3})^2 \Rightarrow 10^{-8} IP > K_{sp}$

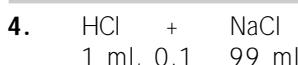
53. $pH = p^{Hln} + \log \frac{In^-}{Hln}$ $6 = 5 + \log \frac{In^-}{Hln}$
 $\frac{In^-}{Hln} = 10$

54. At Half way $[Hln] = In^-$

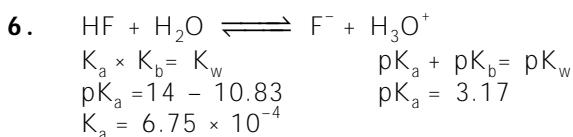
$$pH = 5.5 + \log \frac{[salt]}{[acid]} \quad 5.5 = pK_a + \log \frac{[salt]}{[acid]}$$

$$\log \frac{[salt]}{[acid]} = 0.75 \Rightarrow \frac{[salt]}{[acid]} = 5.62$$

Exercise-02

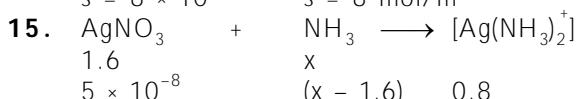


$$[H^+] = \frac{0.1}{100} \Rightarrow 0.001 \quad pH = 3$$



9. $h = \sqrt{\frac{K_h}{C}}$ $h = \sqrt{\frac{K_w}{K_a \times C}}$
 $h = \sqrt{\frac{10^{-14} \times 8.0}{1.3 \times 10^{-9} \times 1}}$ $h = 2.48 \%$

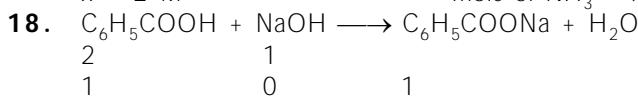
13. $K_{sp} = s^2$ $s = \sqrt{K_{sp}} = \sqrt{6.4 \times 10^{-5}}$
 $s = 8 \times 10^{-3}$ $s = 8 \text{ mol/m}^3$



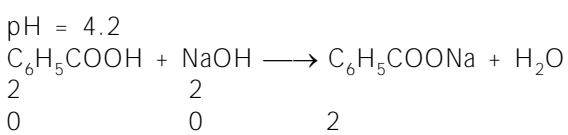
$$K_s = \frac{[Ag(NH_3)_2]^+}{(AgNO_3)(NH_3)}$$

$$10^8 = \frac{0.8}{(5 \times 10^{-8})(x - 1.6)^2} \quad (x - 1.6)^2 = 0.16$$

$$x = 2 \text{ M} \quad \text{mole of } NH_3 = 4$$



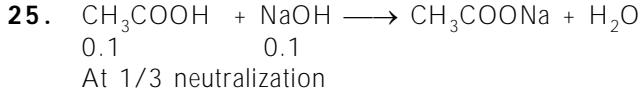
$$pH = pK_a + \log \frac{[salt]}{[acid]} \quad pH = 4.2 + \log \frac{1}{1}$$



$$pH = 7 + \frac{1}{2} pK_a + \frac{1}{2} \log C$$

$$pH = 7 + 2.1 + \frac{1}{2} \log \frac{2}{200}$$

$$pH = 9.1 - 1 \quad pH = 8.1$$



$$2/3 \quad 1/3$$

$$pH = pK_a + \log \frac{(\text{salt})}{(\text{acid})}$$

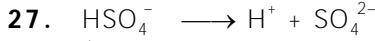
$$pH_1 = pK_a + \log \frac{1/3}{2/3} \quad \dots(1)$$

at 2/3 neutralization

$$pH_2 = pK_a + \log \frac{2/3}{1/3} \quad \dots(2)$$

$$pH_1 - pH_2 = \log \frac{1}{2} - \log 2$$

$$= \log \frac{1}{4} = -2 \log 2$$



$$1 \quad \alpha \quad \alpha$$

$$10^{-2} = \frac{\alpha^2}{1-\alpha} \quad \alpha \Rightarrow 0.09 \quad pH = 1.02$$

Exercise-03

COMPREHENSION BASED QUESTIONS

Comprehension # 1

1. Suppose volume of $HCO_3^- = V \text{ mL}$
millimoles of $HCO_3^- = 5V$
millimoles of $H_2CO_3 = 20$

$$pH = pK_a + \log \frac{[HCO_3^-]}{[H_2CO_3]}$$

$$7.40 = 6.11 + \log \frac{V}{4}, \quad V = 78 \text{ mL}$$

3. If CO_2 escapes, $[H^+]$ decreases, hence pH increases.
(~~if CO_2 escape then $[H^+]$ increases~~)

Comprehension # 2

1. Phosphoric acid with three ionisable hydrogens ions is a tribasic acid. H-atoms are attached to O-atoms,

(Q) If first step is only taken (fn doy ike in dksfy; tk A)

2. If first step is only taken (fn doy ike in dksfy; tk A)

$$\text{pH} = \frac{1}{2}[\text{pK}_{\text{a}_1} - \log c] \quad C = [\text{H}_3\text{PO}_4] = 0.05 \text{ M}$$

$$= \frac{0.05 \times 10}{98} \text{ mol L}^{-1} (\text{M}) = 5.1 \times 10^{-3} \text{ M}$$

$$-\log c = 2.3, \text{ pK}_{\text{a}_1} = 2.12 \text{ pH} = 2.21$$

$$3. \frac{[\text{H}^+]^3 [\text{PO}_4^{3-}]}{[\text{H}_3\text{PO}_4]} = K_{\text{a}_1} K_{\text{a}_2} K_{\text{a}_3}$$

$$3\log [\text{H}^+] + \log [\text{PO}_4^{3-}]$$

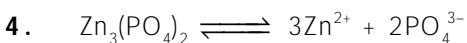
$$= \log K_{\text{a}_1} + \log K_{\text{a}_2} + \log K_{\text{a}_3} - \log [\text{H}_3\text{PO}_4]$$

$$3\text{pH} - \log [\text{PO}_4^{3-}] = \log [\text{H}_3\text{PO}_4] = \text{pK}_{\text{a}_1} + \text{pK}_{\text{a}_2} + \text{pK}_{\text{a}_3}$$

$$21 - \log [\text{PO}_4^{3-}] - 3 = 2.12 + 7.21 + 12.32$$

$$\log [\text{PO}_4^{3-}] = -3.65$$

$$[\text{PO}_4^{3-}] = 2.24 \times 10^{-4} \text{ M}$$

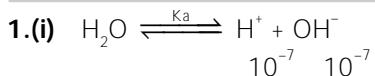


$$\therefore K_{\text{sp}} = [\text{Zn}^{2+}]^3 [\text{PO}_4^{3-}]^2$$

$$9.1 \times 10^{-33} = [\text{Zn}^{2+}]^3 (2.2 \times 10^{-4})^2$$

$$[\text{Zn}^{2+}]^3 = 1.88 \times 10^{-25} [\text{Zn}^{2+}] = 5.73 \times 10^{-9} \text{ M}$$

Exercise-4(A)



$$K_a = \frac{[\text{H}^+][\text{OH}^-]}{[\text{H}_2\text{O}]} = \frac{10^{-7} \times 10^{-7}}{1000/8} = \frac{10^{-14}}{55.5} = 1.8 \times 10^{-16}$$

$$(ii) K_a \times K_b = 10^{-14}$$

$$2. K_a = C\alpha^2 \Rightarrow \frac{\alpha_2}{\alpha_1} = \sqrt{\frac{C_1}{C_2}} = \sqrt{\frac{1}{1/100}} = 10$$

$$3. K_a = C\alpha^2$$

$$\frac{\alpha_1}{\alpha_2} = \sqrt{\frac{K_a_1}{K_a_2}} = \sqrt{\frac{1.8 \times 10^{-5}}{6.2 \times 10^{-10}}}$$

$$4.(a) \text{pH} = \frac{1}{2} \text{pK}_a - \frac{1}{2} \log C$$

$$4.50 \times 2 = \text{pK}_a - \log (0.1)$$

$$9 - 1 = \text{pK}_a \quad \text{pK}_a = 8 \quad K_a = 10^{-8}$$

$$5.(c) [\text{H}^+] = \sqrt{K_a C} = \sqrt{1.8 \times 10^{-6}} = 3 - \log 1.8 = 2.87$$

$$(e) [\text{H}^+] = 10^{-8} + 10^{-7} = 10^{-7} [0.1 + 1]$$

$$\text{pH} = 7 - \log 1.1 = 6.95$$

$$(f) [\text{OH}^-] = 10^{-10} + 10^{-7} = 10^{-7} [1.001]$$

$$\text{POH} = 7 - \log 1.001 = 6.99$$

$$\text{pH} = 7.0004$$

$$(g) [\text{H}^+] = \sqrt{K_a C} = \sqrt{1.8 \times 10^{-5} \times 10^{-6}}$$

$$[\text{H}^+] = \sqrt{1.8 \times 10^{-11}} = \sqrt{18 \times 10^{-12}} = 4.24 \times 10^{-6}$$

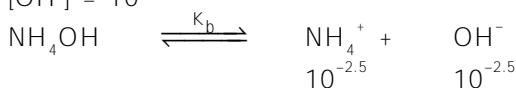
$$\text{pH} = 6 - \log 4.24 = 5.37$$

$$6. \text{pK}_w = 14 - \log 2.56 = 13.59 \approx 13.6$$

$$\text{pH} = \frac{\text{pK}_w}{2} = 6.795$$

$$10. \text{pH} = 11.5 \quad [\text{H}^+] = 10^{-11.5}$$

$$[\text{OH}^-] = 10^{-2.5}$$

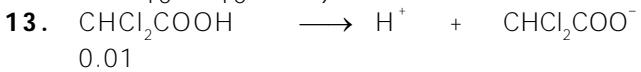


$$10^{-2.5} \quad 10^{-2.5}$$

$$\frac{10^{-5}}{C} = 1.8 \times 10^{-5} \quad C = \frac{1}{1.8} = 0.556 \text{ M}$$

$$11. C = 10^{-2}, [\text{H}^+] = 10^{-3}$$

$$K_a = \frac{10^{-3} \times 10^{-3}}{10^{-2} - 10^{-3}} = \frac{10^{-3}}{9} = 1.1 \times 10^{-4}$$



$$0.01 - x \quad 0.01 + x \quad x$$

$$\frac{x(0.01+x)}{0.01-x} = 2.55 \times 10^{-2}$$

$$0.01x + x^2 = 2.55 \times 10^{-4} - 2.55 \times 10^{-2}x$$

$$x^2 + 0.355x - 0.000255 = 0$$

$$x = \frac{-0.0355 \pm \sqrt{0.04775}}{2} = 1.1 \times 10^{-2}$$

$$\text{CHCl}_2\text{COO}^- = 6.126 \times 10^{-2}$$

14. For weak acid

$$[\text{H}^+] = \sqrt{K_1 C_1 + K_2 C_2 \dots + K_w}$$

$$= \sqrt{1.8 \times 10^{-5} \times 0.02 + 6.4 \times 10^{-5} \times .01 + 10^{-14}}$$

$$= \sqrt{100 \times 10^{-8}}$$

$$[\text{H}^+] = 10^{-3} \quad K_a = \frac{[\text{H}^+][\text{ACO}^-]}{[\text{ACOH}]_2}$$

$$[\text{ACO}^\ominus] = 3.6 \times 10^{-4}$$

$$\text{same } [\text{C}_2\text{H}_5\text{O}_2^\ominus] = 6.4 \times 10^{-4}$$

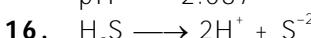
15. HCN is a weak acid so H^+ due to it can neglect As comparision to HF

$$[\text{H}^+] = \sqrt{K C} = \sqrt{6.7 \times 10^{-4} \times 0.1} = \sqrt{67 \times 10^{-6}}$$

$$= 8.18 \times 10^{-3}$$

$$\text{pH} = -\log [8.18 \times 10^{-3}] = 3 - \log [8.18]$$

$$\text{pH} = 2.087$$



$$[\text{H}^+] = 2 \times 10^{-4}, [\text{H}_2\text{S}] = 0.1 \text{ M}$$

$$K = K_1 \times K_2$$

$$K = 10^{-21}$$

$$\Rightarrow 10^{-21} = \frac{[2 \times 10^{-4}]^2 [S^{-2}]}{[0.1]} \quad 10^{-21} = 4 \times 10^{-8} [S^{-2}]$$

$$\frac{1}{4} \times 10^{-14} = [S^{-2}] \quad 2.5 \times 10^{-15} = [S^{-2}]$$

- 17.** (i) $H_3PO_4 \longrightarrow H^+ + H_2PO_4^- \quad K_1 = 7.225 \times 10^{-3}$
 $0.01 \text{ M C} \quad (1-\alpha) \text{ } C\alpha_1 \quad C\alpha_1$
- (ii) $H_2PO_4^- \longrightarrow HPO_4^{2-} + H^+ \quad K_2 = 6.8 \times 10^{-8}$
 $C\alpha_1 (1-\alpha_2) \quad C\alpha_1 \alpha_2 \quad [C\alpha_1]$
- (iii) $HPO_4^{2-} \longrightarrow PO_4^{3-} + H^+ \quad K_3 = 6.8 \times 10^{-8}$
 $\alpha_1 \alpha_2 C_1 (1-\alpha_3) \quad C (\alpha_1 \alpha_2 \alpha_3) \quad [C\alpha_1]$
- $$7.225 \times 10^{-3} = \frac{C\alpha_1^2}{(1-\alpha_1)} = \frac{0.01 \times \alpha_1^2}{1-\alpha}$$

RxN. (i)

$$(1-\alpha) \times 0.7225 = \alpha_1^2$$

$$\alpha_1^2 + 0.7225 \alpha - 0.7225 = 0$$

$$\alpha_1 = 0.562$$

$$\Rightarrow [H^+] = 0.01 \times 0.562 \quad [H^+] = 5.6 \times 10^{-3}$$

$$[H_2PO_4^\ominus] \approx 5.6 \times 10^{-3} \quad \text{RxN. (ii)}$$

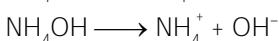
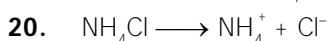
$$6.8 \times 10^{-8} = \frac{[HPO_4^{2-}][H^+]}{[H_2PO_4^-]} \text{ from (i) reaction.}$$

$$\Rightarrow [HPO_4^{2-}] = 6.8 \times 10^{-8} \text{ M} \quad \text{RxN. (iii)}$$

$$4.5 \times 10^{-13} = \frac{[PO_4^{3-}][H^+]}{[HPO_4^{2-}]}$$

$$\frac{4.5 \times 10^{-13} \times 6.8 \times 10^{-8}}{5.6 \times 10^{-3}} = [PO_4^{3-}]$$

$$5.464 \times 10^{-18} = [PO_4^{3-}]$$



$$K_b = \frac{[\text{NH}_4^+][\text{OH}^-]}{[\text{NH}_4\text{OH}]}$$

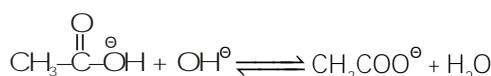
$[\text{NH}_4^+]$ is due to salt because NH_4OH ionise in less amount due to common ions effect

$$1.8 \times 10^{-5} = \frac{0.1 \times [\text{OH}^-]}{0.05} \quad 9 \times 10^{-6} = [\text{OH}^-]$$

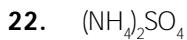


50ml, 0.2M 50ml, 0.1M

10m mol 5m mol



$$\begin{array}{ccccc} 10 & & 5 & & \\ 5 & & 0 & & 5 \\ \text{pH} = \text{pK}_a = 5 - \log 1.8 & & & & \text{pH} = 4.74 \end{array}$$



$$\text{Molarity } (\text{NH}_4)_2\text{SO}_4 = \left(\frac{x}{100} \right)$$

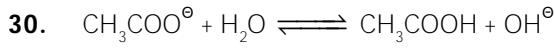
$$\text{Molarity of NH}_4^+ = \left(\frac{2x}{100} \right)$$

$$\text{Molarity of NH}_4\text{OH} = \left(\frac{0.1}{100} \right) = 10^{-3}$$

$$14 - 9.26 = 4.24 + \log \left(\frac{2x/100}{0.1/100} \right)$$

$$0 = \log (20x) \Rightarrow 1 = 20x$$

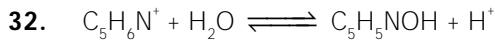
$$x = 1/20 \text{ mole} \quad x = 0.05 \text{ mole}$$



$$0.08 \quad K_b = \frac{10^{-14}}{1.8 \times 10^{-5}}$$

$$\frac{x^2}{0.08} = \frac{10}{1.8} \times 10^{-10} \quad x^2 = \frac{0.8}{1.8} \times 10^{-10}$$

$$x^2 = 0.44 \times 10^{-10} \quad x = 0.66 \times 10^{-5}$$



$$\text{pH} = \frac{1}{2} [\text{pK}_w - \text{pK}_b - \log C]$$

$$2.699 = \frac{1}{2} [14 - \text{pK}_b + 0.6]$$

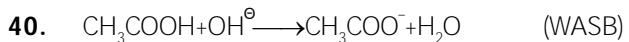
$$2.398 = 14.6 - \text{pK}_b$$

$$\text{pK}_b = 14.6 - 5.398 = 9.802$$

$$K_b = 10^{-9.802}$$

$$\text{38. } \text{pH} = \frac{\text{pK}_1 + \text{pK}_2}{2}$$

$$\text{pH} = \frac{11 + 7 - 2 \log 4.5}{2} = 9 - \log 4.5 = 8.54$$



$$\text{pH} = \frac{1}{2} [\text{pK}_w + \text{pK}_a + \log C]$$

$$= \frac{1}{2} [14 + 5 - \log 1.9 + \log \frac{1}{20}]$$

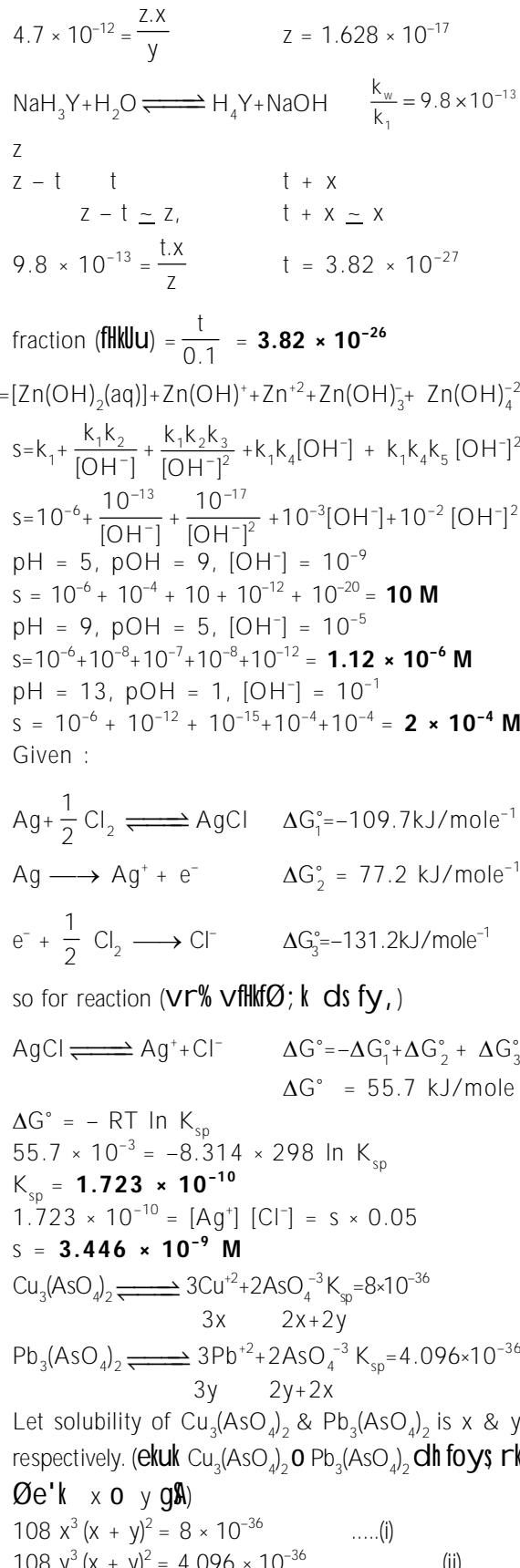
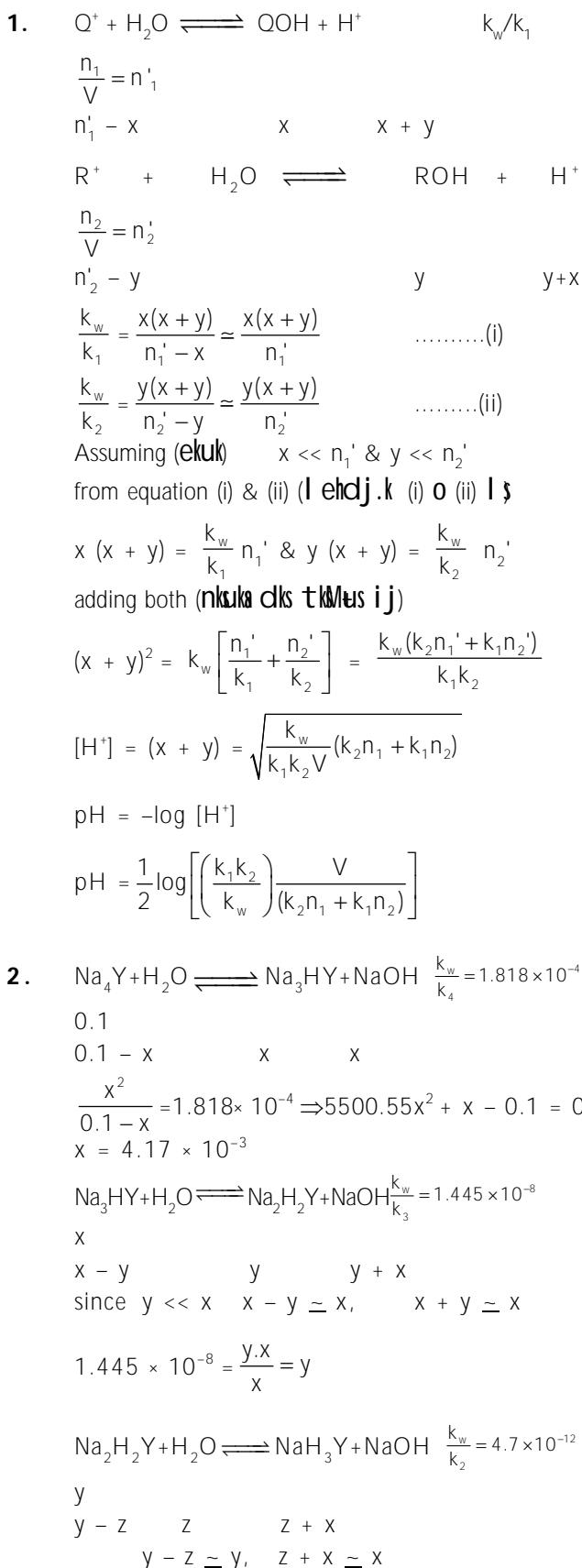
$$= \frac{1}{2} [19 - \log 1.9 - \log 20]$$

$$\text{pH} = \frac{1}{2} [19 - \log 20 \times 1.9] = 8.78$$

$$\text{pOH} = 5.28 \quad [\text{OH}^\ominus] = 10^{-5.28}$$

$$[\text{OH}^\ominus] = 10^{-6} \times 10^{0.72} \quad [\text{OH}^\ominus] = 5.24 \times 10^{-6}$$

Exercise-4(B)



$$\frac{(i)}{(ii)} \Rightarrow \frac{x^3}{y^3} = \frac{8}{4.096} \Rightarrow x = 1.25 y$$

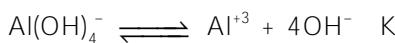
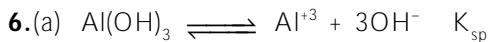
putting this in equation (ii) **(bl sl ehadj.k (ii)ej [kusij])**

$$108 y^3 (2.25 y)^2 = 4.096 \times 10^{-36}$$

$$y = 2.3 \times 10^{-8}$$

$$[Cu^{+2}] = 3x = 8.825 \times 10^{-8}$$

$$[Pb^{+2}] = 3y = 7.1 \times 10^{-8}$$



$$\text{Al(OH)}_3 + \text{OH}^- \rightleftharpoons \text{Al(OH)}_4^- \frac{K_{sp}}{K} = 38.46$$

$$38.46 = \left[\frac{\text{Al(OH)}_4^-}{[\text{OH}^-]} \right] = \frac{10^{-3}}{[\text{OH}^-]}$$

$$[\text{OH}^-] = 2.6 \times 10^{-5}$$

$$\text{pOH} = 4.585$$

$$\text{pH} = 9.415$$

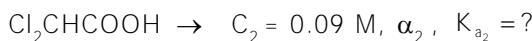
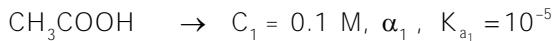
(b) $K_{sp} = [\text{Al}^{+3}] [\text{OH}^-]^3$

$$5 \times 10^{-33} = [1 \times 10^{-3}] [\text{OH}^-]^3$$

$$[\text{OH}^-] = 1.7 \times 10^{-10}$$

$$\text{pOH} = 9.767$$

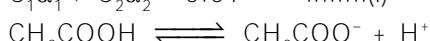
$$\text{pH} = 4.23$$



$$\text{pH} = 1, [\text{H}^+] = 0.1$$

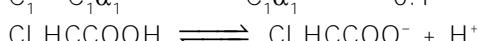
$$0.1 = 0.09 + C_1\alpha_1 + C_2\alpha_2$$

$$C_1\alpha_1 + C_2\alpha_2 = 0.01 \quad \dots\dots(i)$$



$$C_1$$

$$C_1 - C_1\alpha_1 \quad C_1\alpha_1 \quad 0.1$$



$$C_2$$

$$C_2 - C_2\alpha_2 \quad C_2\alpha_2 \quad 0.1$$

$$K_{a_1} = \frac{(C_1\alpha_1)(0.1)}{C_1(1-\alpha_1)} \approx \alpha_1 \times 0.1 = 10^{-5}$$

$$\alpha_1 = 10^{-4}$$

putting this in equation (i) **(bl sl ehadj.k (i)ej [kusij])**

$$10^{-4} \times 0.1 + 0.09\alpha_2 = 0.01$$

$$\alpha_2 = 0.111$$

$$K_{a_2} = \frac{(C_2\alpha_2)(0.1)}{C_2(1-\alpha_2)} = \frac{(0.111)(0.1)}{1-0.111} = 1.248 \times 10^{-2}$$

8. $C = \frac{10 \times 0.935 \times 1000}{17 \times 100} = 5.5 \text{ M}$

$$K_a = C\alpha^2 \Rightarrow 5.5 \times 10^{-6} = 5.5 \alpha^2$$

$$\alpha = 10^{-3}$$

$$[\text{OH}^-] = C\alpha = 5.5 \times 10^{-3}$$

$$\text{pOH} = 2.26, \textbf{pH} = 11.74$$

9. $\ln \frac{k_{w_2}}{k_{w_1}} = \frac{\Delta H}{R} \left(\frac{1}{T_1} - \frac{1}{T_2} \right)$

$$\ln \frac{5.474 \times 10^{-14}}{1.08 \times 10^{-14}} = \frac{\Delta H}{8.314} \left(\frac{1}{298} - \frac{1}{323} \right)$$

$$\Delta H = 51952.6 \text{ J} = 51.95 \text{ kJ/mole}$$

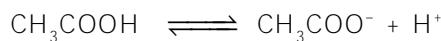
10. In begining $[\text{H}^+] = \sqrt{K_a C}$

$$[\text{H}^+] = \sqrt{1.8 \times 10^{-5}} = 0.004242$$

$$\text{pH} = 2.372$$

On doubling pH, new pH(pH **nqph djustij u; h pH**)
= 4.744

$$[\text{H}^+] = 1.8 \times 10^{-5}$$



$$C$$

$$C - C\alpha \quad C\alpha$$

$$K_a = \frac{(C\alpha)^2}{C(1-\alpha)}, C\alpha = [\text{H}^+] = 1.8 \times 10^{-5}$$

$$1.8 \times 10^{-5} = \frac{(1.8 \times 10^{-5})^2}{C - C\alpha}$$

$$C - C\alpha = 1.8 \times 10^{-5}$$

$$C = 3.6 \times 10^{-5}$$

$$V = \frac{1}{C} = 2.77 \times 10^4 \text{ L}$$

11.(a) $PV = nRT$

$$1 \times 0.959 = n \times 0.0821 \times 298$$

$$n = 0.03919$$

volume of $\text{H}_2\text{O} = 1 \text{ mL}$ (per volume of H_2O)

$(\text{H}_2\text{O} \text{ dk vkl; ru} = 1 \text{ mL} (\text{H}_2\text{O} \text{ dk ifr vkl; ru}))$

$$C = \frac{n}{V} = \frac{0.03919}{10^{-3}} = 39.19 \text{ M}$$

$$pk_b = 3.39 \Rightarrow k_b = 4 \times 10^{-4}$$

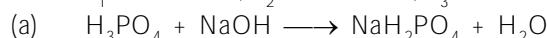
$$[\text{OH}^-] = \sqrt{K_b C} = 0.1252 \text{ M}$$

$$\text{pOH} = 0.9023$$

$$\textbf{pH} = 13.097$$

(b) $M = 0.1252$ for NaOH (NaOH **ds fy, 0.1252**)

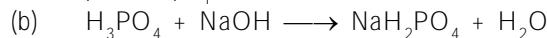
$$k_1 = 7.5 \times 10^{-3}, k_2 = 6.2 \times 10^{-8}, k_3 = 10^{-12}$$



$$6 \quad 3$$

$$3 \quad - \quad 3$$

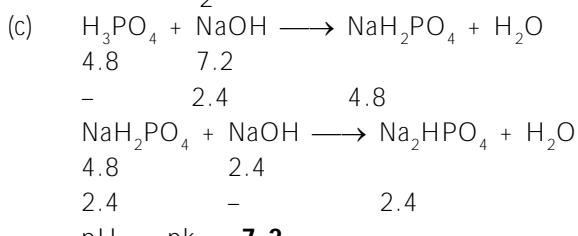
$$\text{pH} = \text{pk}_1 = 2.12$$



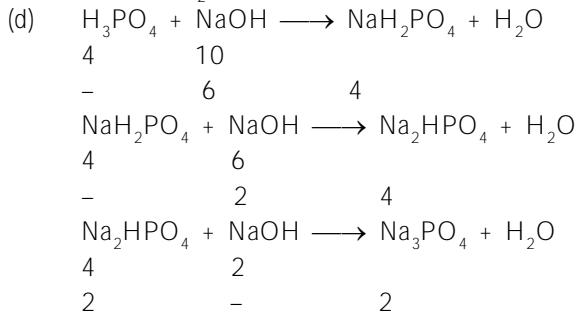
$$6 \quad 6$$

$$- \quad - \quad 6$$

$$\text{pH} = \frac{\text{pk}_1 + \text{pk}_2}{2} = \mathbf{4.66}$$

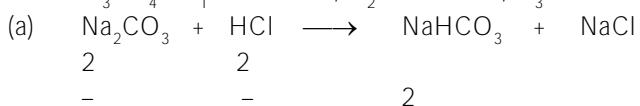


$$\text{pH} = \text{pk}_2 = \mathbf{7.2}$$

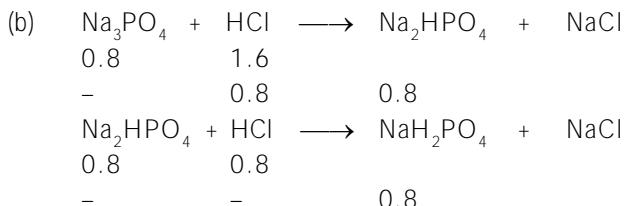


$$\text{pH} = \text{pk}_3 = \mathbf{12}$$

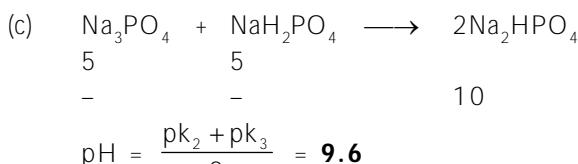
13. For $\text{H}_2\text{CO}_3 \rightarrow \text{k}_1 = 4.2 \times 10^{-7}$, $\text{k}_2 = 4.8 \times 10^{-11}$
 $\text{H}_3\text{PO}_4 \rightarrow \text{k}_1 = 7.5 \times 10^{-3}$, $\text{k}_2 = 6.2 \times 10^{-8}$, $\text{k}_3 = 10^{-12}$



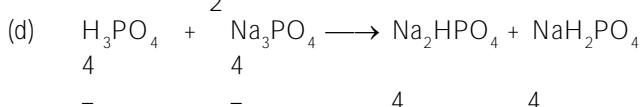
$$\text{pH} = \frac{\text{pk}_1 + \text{pk}_2}{2} = \mathbf{8.347}$$



$$\text{pH} = \frac{\text{pk}_1 + \text{pk}_2}{2} = \mathbf{4.66}$$



$$\text{pH} = \frac{\text{pk}_2 + \text{pk}_3}{2} = \mathbf{9.6}$$



$$\text{pH} = \text{pk}_2 = \mathbf{7.2}$$



At end point m moles of BOH = m moles of HCl (वर्ति बोह डी म नोलेस)

$$0.16 \times V = 4 \quad V = 25 \text{ mL}$$

$$\text{Total volume (दूरी वक्ता रुप)} = 40 + 25 = 65 \text{ mL}$$

$$[\text{BCl}] = \frac{4}{65}$$

since BCl is SAWB

$$\text{pH} = 7 - \frac{1}{2}\text{pk}_b - \frac{1}{2}\log C$$

$$5.23 = 7 - \frac{1}{2}\text{pk}_b - \frac{1}{2}\log \frac{4}{65}$$

$$\text{pk}_b = 4.75$$

Now on further adding NaOH वे इधे NaOH डिस्फेय्कस इज़



$$4 \quad 1.8$$

$$2.2 \quad - \quad 1.8$$

$$\text{pOH} = \text{pk}_b + \log \frac{2.2}{1.8} = 4.837 \Rightarrow \text{pH} = \mathbf{9.1628}$$

15.(a) $\text{pH} = \text{pK}_a + \log \frac{0.06}{0.05}$

$$\text{pH} = 3.744 + \log 1.2 = \mathbf{3.823}$$

(b) On diluting solution 10 times (फॉय; उड़ाक्स 10 गुण रुप डिज़स इज़)

$$[\text{HCOOH}] = 0.005, [\text{HCOONa}] = 0.006$$



$$0.005$$

$$0.005(1-\alpha) \quad 0.005\alpha \quad 0.005\alpha + 0.006$$

$$\text{K}_a = 1.8 \times 10^{-4} = \frac{(0.005\alpha + 0.006)(0.005\alpha)}{0.005(1-\alpha)}$$

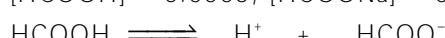
$$\frac{0.005\alpha^2 + 0.006\alpha}{1-\alpha} = 1.8 \times 10^{-4}$$

$$\Rightarrow 27.77\alpha^2 + 34.33\alpha - 1 = 0 \quad \alpha = 0.0285$$

$$[\text{H}^+] = 0.005\alpha = 1.425 \times 10^{-4} \quad \text{pH} = \mathbf{3.846}$$

(c) On further diluting solution by 10 times (फॉय; उड़ाक्स 10 गुण रुप डिज़स इज़)

$$[\text{HCOOH}] = 0.0005, [\text{HCOONa}] = 0.0006$$



$$0.0005$$

$$0.0005(1-\alpha) \quad 0.0005\alpha \quad 0.0005\alpha + 0.0006$$

$$\text{K}_a = 1.8 \times 10^{-4} = \frac{(0.0005\alpha + 0.0006)(0.0005\alpha)}{0.0005(1-\alpha)}$$

$$\frac{0.0005\alpha^2 + 0.0006\alpha}{1-\alpha} = 1.8 \times 10^{-4}$$

$$2.77\alpha^2 + 4.33\alpha - 1 = 0 \Rightarrow \alpha = 0.2047$$

$$[\text{H}^+] = 0.0005\alpha = 1.0235 \times 10^{-4}$$

$$\text{pH} = \mathbf{3.9899}$$

16. Initial (इक्के एक्स) $\text{pOH} = \text{pK}_b = 4.744$

Let x mole of NaOH has been added so (एक्के नाहॉड़ डी से एक्स एक्स फेय्क्स सग्स वर्क)

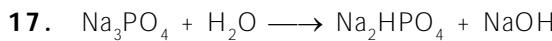
$$[\text{NH}_4^+] = 0.1 + x, [\text{NH}_3] = 0.1 - x$$

$$\text{pOH} = 5.744$$

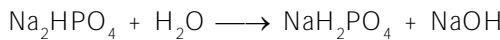
$$5.744 = 4.744 + \log \frac{0.1+x}{0.1-x}$$

$$1 = \log \frac{0.1 + x}{0.1 - x}$$

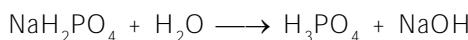
$$\frac{0.1 + x}{0.1 - x} = 10 \Rightarrow x = \frac{0.9}{11} = 0.0818 \text{ moles}$$



$$K = \frac{K_w}{K_3} = 0.0222$$



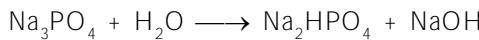
$$K = \frac{K_w}{K_2} = 1.58 \times 10^{-7}$$



$$K = \frac{K_w}{K_3} = 1.4 \times 10^{-12}$$

since equilibrium constant of 2nd & 3rd reaction is very less, $[\text{OH}^-]$ will mainly come from 1st reaction.

(प्र० 2nd o 3rd विक्षेप के लिए कम हैं। इसका उत्तराधिकारी नहीं है)



0.1

0.1-x x x

$$\frac{x^2}{0.1 - x} = 0.0222 \Rightarrow 45x^2 + x - 0.1 = 0$$

$$x = 3.73 \times 10^{-2}$$

$$[\text{OH}^-] = x = 3.73 \times 10^{-2} \text{ M}$$

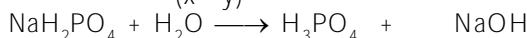


x

$$x - y y y + x$$

$$x - y \leq x, y + x \leq x, \text{ so}$$

$$1.58 \times 10^{-7} = \frac{(y + x)}{(x - y)} y = y$$



y

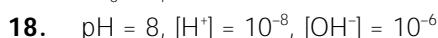
$$y - z z z + x$$

$$y - z \leq y, z + x \leq x$$

$$1.4 \times 10^{-12} = \frac{z(x + z)}{(y - z)} = \frac{z \times x}{y} = \frac{z \times 3.73 \times 10^{-2}}{1.58 \times 10^{-7}}$$

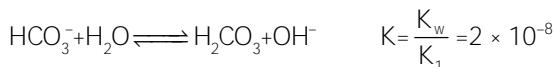
$$z = 5.93 \times 10^{-18}$$

$$[\text{H}_3\text{PO}_4] = z = 5.93 \times 10^{-18} \text{ M}$$



0.0005

$$0.0005 - y - z \quad 10^{-8} \quad y$$



0.0005

$$0.0005 - y - z \quad z \quad 10^{-6}$$

since equilibrium constant for first reaction is very less
(प्र० 1st विक्षेप के लिए कम है। इसका उत्तराधिकारी नहीं है)

$$2 \times 10^{-8} = \frac{z(10^{-6})}{0.0005 - z}$$

$$51z = 0.0005, \Rightarrow z = 9.8 \times 10^{-6}$$

$$[\text{H}_2\text{CO}_3] = 9.8 \times 10^{-6} \text{ M}$$

$$[\text{HCO}_3^-] = 0.0005 - 9.8 \times 10^{-6} = 4.9 \times 10^{-6} \text{ M}$$

$$5 \times 10^{-3} = \frac{10^{-8} \times y}{4.9 \times 10^{-6}}$$

$$[CO_3^{2-}] = y = 2.45 \times 10^{-8} \text{ M}$$

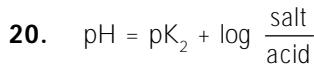


x

$$0.95x \quad 0.05x \quad 0.05x$$

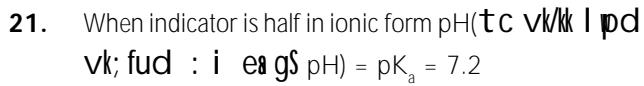
$$6.5 \times 10^{-3} = \frac{(0.05)^2 x}{0.95} \quad x = 2.47$$

$$[\text{H}^+] = 0.05x = 0.1235 \quad \text{pH} = 0.908$$



$$6.7 = 7.2 + \log \frac{y}{0.005}$$

$$y = 1.58 \times 10^{-3} \text{ mole}$$



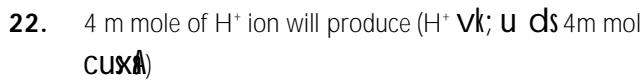
$$\text{pH} = 7.2 + \log 5 = 7.898$$

now with this pH (विक्षेप के लिए pH)

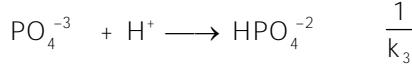
$$7.898 = \text{p}K_{a_1} + \log 4 = \text{p}K_{a_1} = 7.2959$$

again when 50% of new indicator is in ionic form (नया विक्षेप के लिए pH) : i.e. $\text{pH} = 7.2 + \log 5 = 7.898$

$$\text{pH} = \text{p}K_{a_1} = 7.2959$$

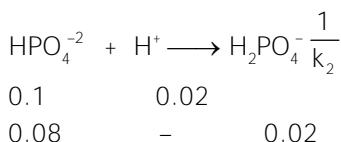


$$[\text{H}^+] = \frac{4 \times 10^{-3}}{0.1} = 0.04$$



$$0.02 \quad 0.04 \quad 0.08$$

$$- \quad 0.02 \quad 0.1$$

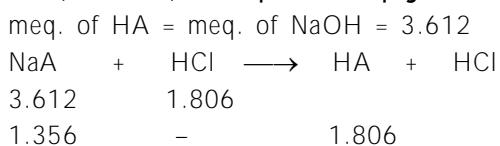


so now they form a buffer solution of HPO_4^{2-} & H_2PO_4^- (vr%vc ; g HPO_4^{2-} o H_2PO_4^- dk cQj foy; u cukrs gA)

$$\text{pH} = \text{pk}_2 + \log \frac{0.08}{0.02} \quad (\text{k}_2 = 6.3 \times 10^{-8})$$

$$\text{pH} = 7.2 + \log 4 = 7.8$$

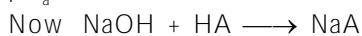
23. At equivalence point (rY; kd fcJnq ij)



$$\text{pH} = \text{pK}_a + \log \frac{[\text{S}]}{[\text{A}]}$$

$$4.92 = \text{pK}_a + \log \frac{1.356}{1.806}$$

$$\text{pK}_a = 5.044$$



$$[\text{NaA}] = \frac{2}{20} = 0.1$$

$$\text{pH} = 7 + \frac{1}{2}\text{pK}_a + \frac{1}{2}\log C = 7 + \frac{5.044}{2} + \frac{1}{2}\log 0.1$$

$$\text{pH} = 9$$

24. In begining let x m mole of BOH are present (i kJEHk

ea ekuk BOH ds x m mole mi fLFkr gA)



x

$$\frac{3x}{4} \quad \frac{x}{4}$$

$$\text{pOH} = \text{pk}_b + \log \frac{1}{3}$$

$$14 - 9.24 = \text{pk}_b - \log 3$$

$$\text{pk}_b = 5.237$$

$$k_b = 5.8 \times 10^{-6}$$



$$\frac{x}{4} = 6, \Rightarrow x = 24$$

$$[\text{BOH}] = \frac{24}{50} = 0.48$$

$$[\text{OH}^-] = \sqrt{k_b \times C} = 1.668 \times 10^{-3}$$

$$\text{pOH} = 2.77$$

$$\text{pH} = 11.22$$

25. (a) pH at one fourth neutralization (, d pkFkkbz mnkl hujdjk i j pH)

$$(\text{pH})_1 = \text{pk}_a + \log \frac{x/4}{3x/4} = \text{pk}_a + \log \frac{1}{3}$$

pH at three fourth neutralization (rhu pkFkkbz mnkl hujdjk i j pH)

$$(\text{pH})_2 = \text{pk}_a + \log \frac{3x/4}{x/4} = \text{pk}_a + \log 3$$

$$\Delta \text{pH} = (\text{pH})_2 - (\text{pH})_1 = 2 \log 3 = 0.9542$$

$$(b) 4.45 = \text{pk}_a + \log \frac{x/3}{2x/3} = \text{pk}_a - \log 2$$

$$\text{pk}_a = 4.751$$

$$(c) \Delta \text{pH} = 2 \quad \text{i.e.}$$

$$(\text{pH})_1 = \text{pk}_a + 1, (\text{pH})_2 = \text{pk}_a - 1$$

$$\text{For } \text{pk}_a + 1 \Rightarrow \frac{[\text{S}]}{[\text{A}]} = 10$$

$$\frac{x}{a-x} = 10 \Rightarrow x = 10a - 10x$$

$$x = \frac{10a}{11}$$

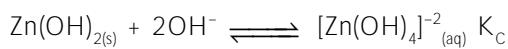
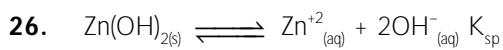
i.e. $\frac{10}{11}$ th stage

$$\text{For } \text{pk}_a - 1 \Rightarrow \frac{[\text{S}]}{[\text{A}]} = \frac{1}{10}$$

$$\frac{x}{a-x} = \frac{1}{10}$$

$$x = \frac{a}{11}$$

i.e. $\frac{1}{11}$ th stage



dissolved Zn(OH)_2 is present in form of Zn^{+2} & $[\text{Zn(OH)}_4]^{2-}$

$$\text{so solubility s} = [\text{Zn}^{+2}] + [\text{Zn(OH)}_4]^{2-}$$

(?kyr Zn(OH)_2 . Zn^{+2} o $[\text{Zn(OH)}_4]^{2-}$ ds: i esmi fLFkr gA)

$$\text{vr% foys rk s} = [\text{Zn}^{+2}] + [\text{Zn(OH)}_4]^{2-}$$

$$s = \frac{K_{sp}}{[\text{OH}^-]^2} + K_C [\text{OH}^-]^2$$

For min. solubility (ഉച്ച ഫോസ്റ്റ് റക്കുഡ്സ്ഫ്യൂഷൻ)

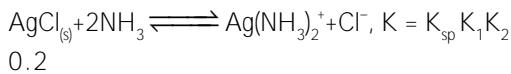
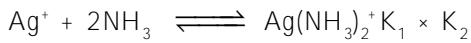
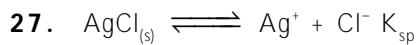
$$\frac{ds}{d[\text{OH}^-]} = 0 \Rightarrow \frac{-2K_{\text{sp}}}{[\text{OH}^-]^3} + 2K_c [\text{OH}^-] = 0$$

$$[\text{OH}^-] = \left(\frac{K_{\text{sp}}}{K_c} \right)^{1/4}$$

$$[\text{OH}^-] = 9.8 \times 10^{-5}$$

$$\text{pOH} = 4.00869$$

$$\text{pH} = 9.9913$$



$$0.2$$

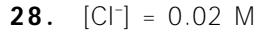
$$0.2 - 2x \quad x \quad x$$

$$K = \frac{x^2}{(0.2 - 2x)} = K_{\text{sp}} K_1 K_2 = 0.002828$$

$$\frac{x}{0.2 - 2x} = 0.05318$$

$$x = 0.009613$$

$$\text{Solubility (ഫോസ്റ്റ് റക്കുഡ്സ്ഫ്യൂഷൻ)} = 9.6 \times 10^{-3} \text{ M}$$



$$0.05$$

$$0.05 - x = 0.05 \quad x \quad 2x$$

$$4 \times 10^{-19} = \frac{x \cdot (2x)^2}{0.05}$$

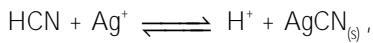
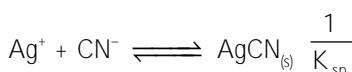
$$\frac{4x^3}{0.05} = 4 \times 10^{-19} \Rightarrow x = 1.7 \times 10^{-7}$$

$$[\text{Ag}] [\text{Cl}] = 1.7 \times 10^{-7} \times 0.02 = 3.4 \times 10^{-9} > K_{\text{sp}}$$

so **AgCl will precipitate.** (വരുത്താനുള്ള ഗ്രാഖാ)

29. After mixing with equal volume (ഉച്ച റൂഡിലുക്കുന്ന ഫേയ്ക്സ്റ്റുസിപ്പർ)

$$[\text{Ag}^+] = 0.01 \text{ M}, \text{ HCN} = 0.01 \text{ M}$$



$$K = \frac{K_a}{K_{\text{sp}}} = 2.25 \times 10^6$$

$$0.01 \quad 0.01$$

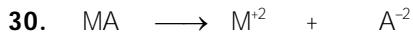
$$x \quad x \quad 0.01$$

since K value is very high almost all of reactant will convert into product

(പിടി കെകു കുറ്റ് മുപ്പ് ഗ്രാഫ്റ്റ് ട്രിക്ക് ഫോഡ് കുടിജി മറിക്കു എം ഫോറ്റ് ഗ്രാഫ്)

$$\frac{0.01}{x^2} = 2.25 \times 10^6 \quad X = 6.6 \times 10^{-5}$$

$$[\text{Ag}^+] = 6.66 \times 10^{-5} \text{ M}$$

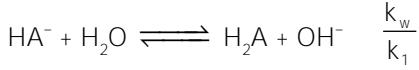


$$s \quad s$$

Let solubility is s. (കുറ്റ് ഫോസ്റ്റ് റക്കുഡ്സ്ഫ്യൂഷൻ)

But some amount of A^{-2} will undergo hydrolysis. Let x is the amount of A^{-2} left in solution.

(യും അംഗീഡിനു കുറ്റ് ട്രിക്ക് വിവരം ഗ്രാഫ് കുറ്റ് ഫോൾ എം ആംഗീഡി അംഗീഡി ഗ്രാഫ്)



$$\frac{k_w}{k_2} = \frac{[\text{HA}^-][\text{OH}^-]}{[\text{A}^{-2}]} \Rightarrow [\text{HA}^-] = \frac{k_w[\text{A}^{-2}]}{k_2[\text{OH}^-]}$$

$$\frac{k_w}{k_1} = \frac{[\text{H}_2\text{A}][\text{OH}^-]}{[\text{HA}^-]} \Rightarrow [\text{H}_2\text{A}] = \frac{k_w[\text{HA}^-]}{k_1[\text{OH}^-]}$$

$$[\text{H}_2\text{A}] = \frac{k_w^2}{k_1 k_2} \frac{[\text{A}^{-2}]}{[\text{OH}^-]^2}$$

From mass balance (20; eku I Urugy I §)

$$S = X + \frac{k_w}{k_2} \frac{[\text{A}^{-2}]}{[\text{OH}^-]} + \frac{k_w^2 [\text{A}^{-2}]}{k_1 k_2 [\text{OH}^-]^2}$$

$$S = X + \frac{[\text{H}^+]X}{k_2} + \frac{[\text{H}^+]^2 X}{k_1 k_2}$$

$$X = \frac{S}{1 + \frac{[\text{H}^+]}{k_2} + \frac{[\text{H}^+]^2}{k_1 k_2}}$$

$$K_{sp} = [\text{M}^{2+}] [\text{A}^{-2}] = S \cdot X = \frac{S^2}{1 + \frac{[\text{H}^+]}{k_2} + \frac{[\text{H}^+]^2}{k_1 k_2}}$$

$$S = \sqrt{K_{sp} \left(1 + \frac{[\text{H}^+]}{k_2} + \frac{[\text{H}^+]^2}{k_1 k_2} \right)}$$

1. $\because \text{pH} = 1 ; \text{H}^+ = 10^{-1} = 0.1 \text{ M}$
 $\text{pH} = 2 ; \text{H}^+ = 10^{-2} = 0.01 \text{ M}$
 $\therefore M_1 = 0.1 \quad V_1 = 1$
 $M_2 = 0.01 \quad V_2 = ?$

From

$$\begin{aligned} M_1 V_1 &= M_2 V_2 \\ 0.1 \times 1 &= 0.01 \times V_2 \\ V_2 &= 10 \text{ litre} \end{aligned}$$

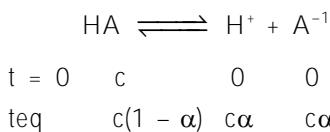
\therefore volume of water added = $10 - 1 = 9$ litre.

2. $\text{H}^+ = C\alpha ; \alpha = \frac{[\text{H}^+]}{C} \quad \text{or} \quad \alpha = \frac{10^{-3}}{0.1} = 10^{-2}$
 $K_a = C \alpha^2 = 0.1 \times 10^{-2} \times 10^{-2} = 10^{-5}$
3. $\text{Cr(OH)}_3(s) \rightleftharpoons \text{Cr}^{3+}(\text{aq.}) + 3\text{OH}^-(\text{aq.})$
 $27S^4 = K_{sp}$

$$S = \left(\frac{K_{sp}}{27} \right)^{1/4} = \left(\frac{1.6 \times 10^{-30}}{27} \right)^{1/4}$$

4. $\text{pH} = 5$ means

$$[\text{H}^+] = 10^{-5}$$



$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]} = \frac{(c\alpha)^2}{c(1-\alpha)} = \frac{[\text{H}^+]^2}{c - [\text{H}^+]}$$

But, $[\text{H}^+] \ll C$

$$\therefore K_a = (10^{-5})^2 = 10^{-10}$$

5. $\text{AgBr} \rightleftharpoons \text{Ag}^+ + \text{Br}^-$

$$K_{sp} = [\text{Ag}^+][\text{Br}^-]$$

For precipitation to occur

Ionic product $>$ Solubility product

$$[\text{Br}^-] = \frac{K_{sp}}{[\text{Ag}^+]} = \frac{5 \times 10^{-13}}{0.05} = 10^{-11}$$

i.e., precipitation just starts when 10^{-11} moles of KBr is added to 1ℓ AgNO₃ solution

\therefore Number of moles of Br⁻ needed from KBr = 10^{-11}

$$\therefore \text{Mass of KBr} = 10^{-11} \times 120 = 1.2 \times 10^{-9} \text{ g}$$

6. $\text{H}_2\text{CO}_3(\text{aq}) + \text{H}_2\text{O}(\ell) \rightleftharpoons \text{HCO}_3^-(\text{aq}) + \text{H}_3\text{O}^+(\text{aq})$
 $0.034-x \quad x \quad x$

$$K_1 = \frac{[\text{HCO}_3^-][\text{H}_3\text{O}^+]}{[\text{H}_2\text{CO}_3]} = \frac{x \times x}{0.034 - x}$$

$$\Rightarrow 4.2 \times 10^{-7} = \frac{x^2}{0.034} \Rightarrow x = 1.195 \times 10^{-4}$$

As H₂CO₃ is a weak acid so the concentration of H₂CO₃ will remain 0.034 as $0.034 \gg x$.

$$x = [\text{H}^+] = [\text{HCO}_3^-] = 1.195 \times 10^{-4}$$

$$\begin{array}{ccccc} \text{Now, } \text{HCO}_3^-(\text{aq}) + \text{H}_2\text{O}(\ell) & \rightleftharpoons & \text{CO}_3^{2-}(\text{aq}) & + & \text{H}_3\text{O}^+(\text{aq}) \\ x-y & & y & & y \end{array}$$

As HCO₃⁻ is again a weak acid (weaker than H₂CO₃) with $x \gg y$.

$$K_2 = \frac{[CO_3^{2-}][H_3O^+]}{[HCO_3^-]} = \frac{y \times (x+y)}{(x-y)}$$

Note : $[H_3O^+] = H^+$ from first step(x) and from second step(y) = $(x + y)$

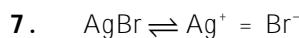
[As $x \gg y$ so $x + y \approx x$ and $x - y \approx x$]

$$\text{So, } K_2 = \frac{y \times x}{x} = y$$

$$\Rightarrow K_2 = 4.8 \times 10^{-11} = y = [CO_3^{2-}]$$

So the concentration of $[H^+] \approx [HCO_3^-]$ = concentrations obtained from the first step. As the first step. As the dissociation will be very low in second step so there will be no change in these concentrations.

$$[H^+] = [HCO_3^-] = 1.195 \times 10^{-4} \text{ & } [CO_3^{2-}] = 4.8 \times 10^{-11}$$



$$K_{sp} = [\text{Ag}^+] [\text{Br}^-]$$

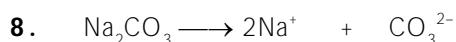
For precipitation to occur Ionic product > Solubility product

$$[\text{Br}^-] = \frac{K_{sp}}{[\text{Ag}^+]} = \frac{5 \times 10^{-13}}{0.05} = 10^{-11}$$

i.e., precipitation just starts when 10^{-11} moles of KBr is added to 1ℓ AgNO₃ solution

$$\therefore \text{Number of moles of Br}^- \text{ needed from KBr} = 10^{-11}$$

$$\therefore \text{Mass of KBr} = 10^{-11} \times 120 = 1.2 \times 10^{-9} \text{ g}$$

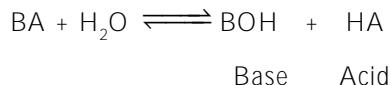


$$1 \times 10^{-4} \text{ M} \quad 1 \times 10^{-4} \text{ M} \quad 1 \times 10^{-4} \text{ M}$$

$$K_{sp}(\text{BaCO}_3) = [\text{Ba}^{2+}] [\text{CO}_3^{2-}]$$

$$[\text{Ba}^{2+}] = \frac{5.1 \times 10^{-9}}{1 \times 10^{-4}} = 5.1 \times 10^{-5} \text{ M}$$

9. It corresponds to choice (c) which is correct answer.



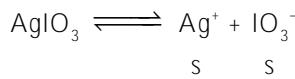
Now pH is given by

$$\text{pH} = \frac{1}{2} \text{pK}_w + \frac{1}{2} \text{pK}_a - \frac{1}{2} \text{pK}_b$$

Substituting given values, we get

$$\text{pH} = \frac{1}{2} (14 + 4.80 - 4.78) = 7.01$$

10. Let s = solubility



$$K_{sp} = [\text{Ag}^+] [\text{IO}_3^-] = s \times s = s^2$$

$$\text{Given } K_{sp} = 1 \times 10^{-8}$$

$$\therefore s = \sqrt{K_{sp}} = \sqrt{1 \times 10^{-8}}$$

$$= 1.0 \times 10^{-4} \text{ mol/lit}$$

$$= 1.0 \times 10^{-4} \times 283 \text{ g/lit}$$

(∴ Molecular mass of Ag IO₃ = 283)

$$= \frac{1.0 \times 10^{-4} \times 283 \times 100}{1000} \text{ gm/100ml}$$

$$= 2.83 \times 10^{-3} \text{ gm/100 ml}$$



$$S \quad 2s$$

Where s is the solubility of MX₂

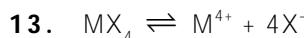
$$\text{then } K_{sp} = 4s^3 ; (2s)^2 = 4 \times 10^{-12} = 4s^3 ; s = 1 \times 10^{-4}$$

$$\therefore [\text{M}^{++}] = s = 1 \times 10^{-4}$$

12. $\text{pH} = -\log[\text{H}^+] = \log \frac{1}{[\text{H}^+]}$

$$5.4 = \log \frac{1}{[\text{H}^+]}$$

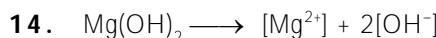
$$\text{On solving, } [\text{H}^+] = 3.98 \times 10^{-6}$$



$$S \quad 4s$$

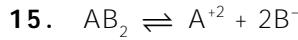
$$K_{sp} = [s] [4s]^4 = 256 s^5$$

$$\therefore s = \left(\frac{K_{sp}}{256} \right)^{1/5}$$



$$x \quad 2x$$

$$K_{sp} = [\text{Mg}] [\text{OH}]^2 = [x] [2x]^2 = x \cdot 4x^2 = 4x^3$$



$$[\text{A}] = 1.0 \times 10^{-5}, [\text{B}] = [2.0 \times 10^{-5}],$$

$$K_{sp} = [B]^2[A] = [2 \times 10^{-5}]^2 [1.0 \times 10^{-5}] = 4 \times 10^{-15}$$