

TOPICS : Equilibrium I and II

- In which one of the following reactions, K_p is less than K_c ?
 - $2\text{SO}_{3(g)} \rightleftharpoons 2\text{SO}_{2(g)} + \text{O}_{2(g)}$
 - $\text{N}_{2(g)} + 3\text{H}_{2(g)} \rightleftharpoons 2\text{NH}_{3(g)}$
 - $\text{PCl}_{5(g)} \rightleftharpoons \text{PCl}_{3(g)} + \text{Cl}_{2(g)}$
 - $\text{H}_{2(g)} + \text{I}_{2(g)} \rightleftharpoons 2\text{HI}_{(g)}$
- The equilibrium constants for the reaction,

$$\text{Zn}_{(s)} + \text{Cu}^{2+}_{(aq)} \rightleftharpoons \text{Zn}^{2+}_{(aq)} + \text{Cu}_{(s)}$$
 and

$$\text{Cu}_{(s)} + 2\text{Ag}^{+}_{(aq)} \rightleftharpoons \text{Cu}^{2+}_{(aq)} + 2\text{Ag}_{(s)}$$
 are K_1 and K_2 respectively. Then the equilibrium constant for the reaction,

$$\text{Zn}_{(s)} + 2\text{Ag}^{+}_{(aq)} \rightleftharpoons \text{Zn}^{2+}_{(aq)} + 2\text{Ag}_{(s)}$$
 will be
 - $K_1 + K_2$
 - $K_1 \times K_2$
 - K_1/K_2
 - $K_1 - K_2$
- For the system $\text{Ag}_{(g)} + 2\text{B}_{(g)} \rightleftharpoons \text{C}_{(g)}$, the equilibrium concentrations are

$$A = 0.06 \text{ mol L}^{-1}, B = 0.12 \text{ mol L}^{-1}, C = 0.216 \text{ mol L}^{-1}$$
 The K_{eq} for the reactions is
 - 250
 - 416
 - 4×10^{-3}
 - 125
- If 0.2 mol of $\text{H}_{2(g)}$ and 2.0 mol of $\text{S}_{(s)}$ are mixed in a 1 dm^3 vessel at 90°C , the partial pressure of $\text{H}_2\text{S}_{(g)}$ formed according to the reaction

$$\text{H}_{2(g)} + \text{S}_{(s)} \rightleftharpoons \text{H}_2\text{S}_{(g)}, (K_p = 6.8 \times 10^{-2})$$
 would be
 - 0.19 atm
 - 0.38 atm
 - 0.6 atm
 - 0.072 atm
- Which one of the following mixtures will give a solution with pH greater than 7 ?
 - 50 mL of 0.1 M HCL + 50 mL of 0.2 M NaCl
 - 50 mL of 0.1 M HCL + 50 mL of 0.1 M CH_3COONa
 - 50 mL of 0.1 M CH_3COOH + 50 mL of 0.1 M NaOH
 - 50 mL of 0.2 M H_2SO_4 + 50 mL of 0.3 M NaOH
- 40 mL of 0.1 M ammonium hydroxide is mixed with 20 mL of 0.1 M HCL. What is the pH of the mixture? ($\text{p}K_b$ of ammonia solution is 4.74)
 - 4.74
 - 2.26
 - 9.26
 - 5
- What is the decreasing order strength of the bases

$$\text{OH}^-, \text{NH}_2\text{H}-\text{C}\equiv\text{C}^- \text{ and } \text{C}\equiv\text{C}^- > \text{OH}^-$$
 - $\text{CH}_3\text{CH}_2^- > \text{NH}_2^- > \text{H}-\text{C}\equiv\text{C}^- > \text{OH}^-$
 - $\text{H}-\text{C}\equiv\text{C}^- < \text{CH}_3-\text{CH}_2^- > \text{NH}_2^- > \text{OH}^-$
 - $\text{OH}^- > \text{NH}_2^- > \text{H}-\text{C}\equiv\text{C}^- > \text{CH}_3\text{CH}_2^-$
 - $\text{NH}_2^- > \text{H}-\text{C}\equiv\text{C}^- > \text{OH}^- > \text{CH}_3\text{CH}_2^-$
- If S_1, S_2, S_3 and S_4 , are the solubilities of AgCl in water, in 0.01 M CaCl_2 , in 0.01 M NaCl and in 0.05 M AgNO_3 respectively at a certain temperature, the correct order of solubilities is
 - $S_1 > S_2 > S_3 > S_4$
 - $S_1 > S_3 > S_2 > S_4$
 - $S_1 > S_2 = S_3 > S_4$
 - $S_1 > S_3 > S_4 > S_2$
- The pH of an aqueous solution of CH_3COONa of concentrated C (M) is given by
 - $7 - \frac{1}{2}\text{p}K_a + \frac{1}{2}\log C$
 - $\frac{1}{2}\text{p}K_w + \frac{1}{2}\text{p}K_b + \frac{1}{2}\log C$
 - $\frac{1}{2}\text{p}K_w - \frac{1}{2}\text{p}K_b - \frac{1}{2}\log C$
 - $\frac{1}{2}\text{p}K_w + \frac{1}{2}\text{p}K_b + \frac{1}{2}\log C$
- A buffer solution contains 0.1 mole of sodium acetate dissolved in 1000 cm^3 of 0.1 M acetic acid. To the above buffer solution, 0.1 mole sodium acetate is further added and dissolved. The pH of the resulting
 - $\text{p}K_a$
 - $\text{p}K_a + 2$
 - $\text{p}K_a - \log 2$
 - $\text{p}K_a + \log 2$

TOPICS : Equilibrium I and II (SOLUTION)

1. (b) : $K_p = K_c(RT)^{\Delta n_g}$, $K_p < K_c$ when Δn_g is negative.
2. (b) : $K =$ product of equilibrium constants of step reactions.

3. (a) : $K = \frac{[C]}{[A][B]^2} = \frac{0.216}{0.06 \times (0.12)^2} = 250$

4. (b) : Suppose x moles of H_2S have formed, then at equilibrium, $[H_2] = (0.2 - x)$, $[H_2S] = x$

$$p_{H_2} = \frac{0.2 - x}{0.2 - x + x} = \frac{0.2 - x}{0.2} \times P$$

$$p_{H_2S} = \frac{x}{0.2 - x + x} = \frac{x}{0.2} \times P$$

$$K_p = \frac{p_{H_2S}}{p_{H_2}} \text{ i.e., } 6.8 \times 10^{-2} = \frac{x}{0.2 - x}$$

$$\text{or } 0.068(0.2 - x) = x \text{ or } x = 0.0127 \text{ mol}$$

Pressure of 0.0127 mol of H_2S at 363 K in 1 L vessel,

$$P = \frac{nRT}{V} = \frac{0.0127 \times 0.0821 \times 363}{1} = 0.38 \text{ atm}$$

5. (c) : All except (c) will be acidic.
6. (c) : Since, ammonium hydroxide is 50% neutralised, hence $[\text{Salt}] = [\text{Base}]$
- $$pOH = pK_a + \log_{10} \frac{[\text{salt}]}{[\text{base}]} = 4.74 + \log_{10} 1$$
- $$pOH = 4.74$$
- $$\therefore pH = 14 - 4.74 = 9.26$$
7. (a) : The conjugate acids of the given bases are H_2O , NH_3 , C_2H_2 , CH_3CH_3 . Their acidic strength are in the order $H_2O > C_2H_2 > NH_3 > CH_3CH_3$. A weak acid has a strong conjugate base.

8. (b) : $0.01 \text{ M CaCl}_2 \equiv 0.02 \text{ M Cl}^-$,
 $0.01 \text{ M NaCl} \equiv 0.01 \text{ M Cl}^-$,
 $0.05 \text{ M AgNO}_3 \equiv 0.05 \text{ M Ag}^+$
 $[\text{Ag}^+][\text{Cl}^-] = K_{sp} \text{ (const.)}$.

The concentration of common ion $\propto \frac{1}{\text{solubility}}$

9. (d) : pH of hydrolysed salt of strong base and weak acid is given by

$$\text{pH} = \frac{1}{2} \text{p}K_w + \frac{1}{2} \text{p}K_a + \frac{1}{2} \log C$$

10. (d) : Number of moles of acetic acid = 0.1 mole
Total number of moles of sodium acetate present in buffer solution = $0.1 + 0.1 = 0.2$ mole

$$\begin{aligned} \therefore \text{pH} &= \text{p}K_a + \log \frac{[\text{CH}_3\text{COONa}]}{[\text{CH}_3\text{COOH}]} \\ &= \text{p}K_a + \log \frac{0.2}{0.1} = \text{p}K_a + \log 2 \\ \therefore \text{pH} &= \text{p}K_a + \log 2 \end{aligned}$$