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# CHEMISTRY

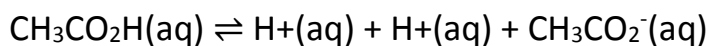
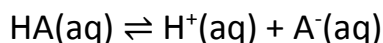
## PHYSICAL CHEMISTRY

Level & Board	CIE (A-LEVEL)
TOPIC:	EQUILIBRIA
PAPER TYPE:	SOLUTION - 1
TOTAL QUESTIONS	10
TOTAL MARKS	75

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**Equilibria - 1**

1)

(a) A weak acid dissociates incompletely in water to give  $H^+$ (b) (i)  $C_i / \text{mold m}^{-3}$       0.100      -      - $C_{eqm} / \text{mold m}^{-3}$     0.100 - x      x      x

$$K_C \frac{[H^+][CH_3CO_2^-]}{[CH_3CO_2H]}$$

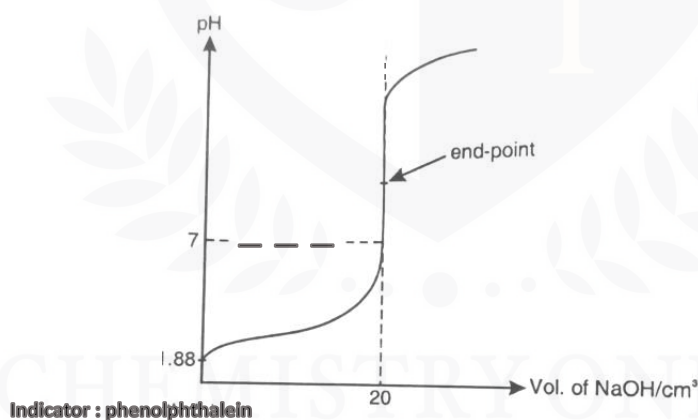
$$1.74 \times 10^{-5} = \frac{x^2}{0.100 - x}$$

Assume  $x \ll 0.100$  since  $CH_3CO_2H$  is a weak acid.

$$1.74 \times 10^{-5} = \frac{x^2}{0.100}$$

$$x = 1.319 \times 10^{-2} \text{ mol dm}^{-3}$$

$$pH = -\lg[H^+] = 1.88$$



$$(c) n_{HCl} = 0.200 \times \frac{14}{1000} = 2.8 \times 10^{-3} \text{ mol}$$

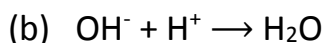
$$n_N = 2.8 \times 10^{-3} \text{ mol}$$

$$m_N = 2.8 \times 10^{-3} \times 14 = 3.92 \times 10^{-2} \text{ g}$$

$$\% N = \frac{3.92 \times 10^{-2}}{0.100} \times 100 = 39.2\%$$

2)

(a) A buffer is usually a mixture of a weak acid and its salt or a weak base and its salt, and it is able to resist pH changes when a little acid or base is added to it.



When  $\text{OH}^-$  it is removed by reacting with the  $\text{H}^+$  in the system.

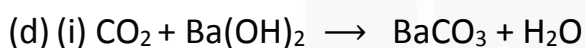
Hence, the pH does not increase.

(c)  $K_a \frac{[\text{H}^+][\text{HCO}_3^-]}{[\text{CO}_2]}$

$$7.90 \times 10^{-7} = [\text{H}^+] \cdot \frac{20}{1}$$

$$[\text{H}^+] = 3.95 \times 10^{-8} \text{ mol dm}^{-3}$$

$$\therefore \text{pH} = -\lg[\text{H}^+] = 7.40$$



(ii) Mr of  $\text{BaCO}_3 = 137 + 12 + 3 \times 16 = 197$

$$n_{\text{BaCO}_3} = \frac{0.600}{197} = 3.05 \times 10^{-3} \text{ mol}$$

Hence, total amount of  $\text{CO}_2 + \text{HCO}_3^- = 3.05 \times 10^{-3} \text{ mol}$ .

$$\frac{[\text{HCO}_3^-]}{[\text{CO}_2]} = \frac{20}{1}$$

$$[\text{HCO}_3^-] = 20[\text{CO}_2]$$

$$[\text{HCO}_3^-] + [\text{CO}_2] = 21[\text{CO}_2]$$

$$\frac{3.05 \times 10^{-3}}{\frac{100}{100}} = 21[\text{CO}_2]$$

$$[\text{CO}_2] = 1.45 \times 10^{-3} \text{ mol dm}^{-3}$$

3)

(b) (i)  $K_{sp} = [\text{Sr}^{2+}][\text{IO}_3^-]^2 \text{ mol}^3 \text{ dm}^{-9}$

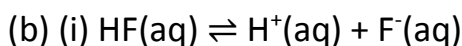
(ii) Let  $[\text{Sr}^{2+}] = x$ .

Then  $[\text{IO}_3^-] = 2x$

$$x(2x)^2 = 1.1 \times 10^{-9}$$

$$x = 6.50 \times 10^{-4} \text{ mol dm}^{-3}$$

4)



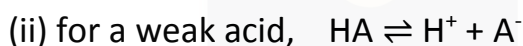
$$[\text{H}^+] = 0.08 \times 0.1$$

$$= 8 \times 10^{-3} \text{ mol dm}^{-3}$$

$$\text{pH} = -\lg[\text{H}^+] = 2.1$$

(ii) The dissociation of HF involves breaking the H – F bond. This process is highly endothermic as indicated by the high bond energy.

5)



$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]}$$

$$(iii) K_a = 10^{-2.57}$$

$$= 2.69 \times 10^{-3} \text{ mol dm}^{-3}$$



(ii) The presence of electronegative F and Cl withdraws negative charge away from  $-\text{CO}_2^-$  by inductive effect. This spreads the charge and stabilizes the anion. Hence, the acids  $\text{FCH}_2\text{CO}_2\text{H}$  and  $\text{ClCH}_2\text{CO}_2\text{H}$  are stronger. F is more electronegative than Cl. Hence, the stabilizing effect is greater and  $\text{FCH}_2\text{CO}_2\text{H}$  is more acidic than  $\text{ClCH}_2\text{CO}_2\text{H}$ .

6)

(b) (i)  $K_p = \frac{P_{\text{N}_2\text{O}_3}^2}{P_{\text{N}_2\text{O}_5} P_{\text{NO}}^2}$

(ii)  $K_c = \frac{0.25^2}{0.75 \times 1.5^2} = 0.0370 \text{ atm}^{-1}$

(c) (i) According to Le Chatelier's principle, when temperature increases, the position of equilibrium shifts to the left whereby the excess heat is removed

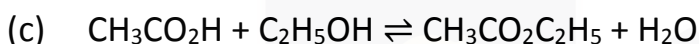
since the backward reaction is endothermic.

- (ii) When pressure is increased, according to Le Chatelier's principle, the position of equilibrium shifts to the right where there is a reduction in the number of gaseous particles. This helps to reduce the increased pressure.

7)

(a) The rate of forward reaction equals the rate of backward reaction.

$$(b) K_a = \frac{[CH_3CO_2C_2H_5][H_2O]}{[CH_3CO_2H][C_2H_5OH]}$$



Initial moles

Let x be the number of moles of reactants used and that of products produced

equilibrium moles (0.5 - x) (0.5 - x) (0.1 + x) (0.1 + x)

equilibrium

concentration /  $\frac{(0.5-x)}{V}$   $\frac{(0.5-x)}{V}$   $\frac{(0.1-x)}{V}$   $\frac{(0.1-x)}{V}$

mole dm<sup>-3</sup>

$$K_c = \frac{\left(\frac{(0.1-x)}{V}\right)\left(\frac{(0.1-x)}{V}\right)}{\left(\frac{(0.5-x)}{V}\right)\left(\frac{(0.5-x)}{V}\right)}$$

$$4 = \frac{(0.1+x)^2}{(0.5-x)^2}$$

$$(0.5 - x)^2 4 = (0.1 + x)^2$$

$$2(0.5 - x) = (0.1 + x)$$

$$1 - 2x = 0.1 + x$$

$$x = 0.3$$

$$\therefore n(CH_3CO_2H) = n(C_2H_5OH) = 0.5 - 0.3 = 0.2$$

$$n(CH_3CO_2C_2H_5) = n(H_2O) = 0.1 + 0.3 = 0.4$$

alcohol Reagent(s) and conditions	$\text{CH}_3\text{CH}_2\text{CH}_2\text{OH}$	$\text{CH}_3\text{CH}_2\text{CH}(\text{OH})\text{CH}_3$	$(\text{CH}_3)_3\text{COH}$
Red phosphorus and iodine heat under reflux		$\text{CH}_3\text{CH}_2\text{CHCH}_3$ 	
Concentrated $\text{H}_2\text{SO}_4$ heat			$\text{CH}_2-\text{C}=\text{CH}_2$   CH
$\text{Cr}_2\text{O}_7^{2-}/\text{H}^+$ Heat under reflux	$\text{CH}_3\text{CH}_2\text{CH}_3\text{CO}$	$\text{CH}_3\text{CH}_2\text{COOH}_3$	no reaction

8)

$$(i) K_a = \frac{[\text{RCO}_2^-][\text{H}^+]}{[\text{RCO}_2\text{H}]}$$

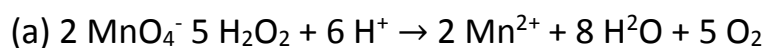
$$(i) \text{p}K_a = -\log_{10} K_a$$

(b) (i) Acid strength increases down the table. As the number of chlorine atoms increase, electron withdrawing effect increases making anion more stable.

(ii) Chlorine atom is further away from O – H in number 4, so it has less influence upon – OH, making acid to be weaker than in number 2.

$$\begin{aligned} (iii) \text{pH} &= \frac{1}{2}(\text{p}K_a - \log_{10}[\text{Acid}]) \\ &= \frac{1}{2}(4.9 - \log_{10}0.01) \\ &= \frac{1}{2}(4.9 + 2) = 3.45 \approx 3.5 \end{aligned}$$

9)



$$(b) E_{\text{cell}}^\ominus = 1.52 - 0.68 = + 3 \times 0.84 \text{ V}$$

(c) (i) Colour changes from purple to colourless

Since  $\text{H}_2\text{O}_2 : \text{MnO}_4^- = 5 : 2$

$$n(\text{H}_2\text{O}_2) = \frac{5}{2} \times 3 \times 10^{-4} \text{ in } 25 \text{ cm}^3$$

$$\therefore [\text{H}_2\text{O}_2] = 7.5 \times \frac{1000}{25} = 3.0 \times 10^{-2} \text{ mol dm}^{-3}$$

10)

(a) High temperature and pressure provide enough energy to break  $\text{N} \equiv \text{N}$  bond.

(b) (i)  $\text{C}_1$   $\text{CO}_1$  hydrocarbons and  $\text{SO}_2$  and  $\text{H}_2\text{S}$   $\text{NO}_x$

(ii) Pt or Pd and Rh

(iii)  $2\text{NO} + 2\text{CO} \rightarrow 2\text{CO}_2 + \text{N}_2$  or  $2\text{NO} + \text{C} \rightarrow \text{CO}_2 + \text{N}_2$

(c) (i)  $K_c = \frac{[\text{NO}]^2[\text{Cl}_2]}{[\text{NOCl}]^2}$ , unit:  $\text{mol dm}^{-3}$

$$(ii) 230^\circ\text{C}: K_c = \frac{(1.46 \times 10^{-2})^2 \times (1.15 \times 10^{-2})}{(2.33 \times 10^{-3})^2} = 4.5 \times 10^{-3} \text{ mol dm}^{-3}$$

$$465^\circ\text{C}: K_c = \frac{(7.63 \times 10^{-3})^2 \times (2.14 \times 10^{-4})}{(3.68 \times 10^{-4})^2} = 9.2 \times 10^{-2} \text{ mol dm}^{-3}$$

(iii) Endothermic because  $K_c$  increases with temperature.

(d) (i) Equilibrium moves to R.H.S. because more moles on R.H.S.

(ii) No change to equilibrium position.

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I am Sorry !!!!!



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- Founder & CEO of Chemistry Online Tuition Ltd.
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