

## CHEM1011 WORKSHEET B

### SAMPLE EXAM QUESTIONS

#### ACIDS, BASES AND pH

##### Question 1.

(a) Complete the following table of conjugate acid/base pairs.

Conjugate acids have one more 'H' than the conjugate base!

Acid	Base
HF	$F^-$
$H_3PO_4$	$H_2PO_4^-$
$H_2O$	$OH^-$
$HNO_2$	$NO_2^-$
$CO_3^{2-}$	— (there isn't one)

(b) Calculate the pH of a 0.15 M solution of vitamin B<sub>5</sub> (pantothenic acid) given  $K_a = 3.95 \times 10^{-5}$ .

$$K_a = \frac{x^2}{[HA]} \text{ where } x = [H^+]$$

$$\text{rearrange: } x = (3.95 \times 10^{-5} \times 0.15)^{1/2}$$

$$\text{so: } pH = -\log [H^+] = 2.61 \text{ (answer to 2 significant figures)}$$

Check: the least accurate value determines the number of significant figures!

(c) A buffer is made up to contain  $[NH_4^+] = 0.2500$  M and  $[NH_3] = 0.1234$  M. Given that  $pK_a(NH_4^+) = 9.24$ , calculate the pH of the buffer.

Use the Henderson-Hasselbalch equation

$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

$$= 9.24 + \log (0.1234/0.2500) = 8.93 \text{ (answer to 2 significant figures)}$$

Check: if more of acid than base, then pH is lower than  $pK_a$ !

**Question 2.**

(a) Calculate the pH of an aqueous buffer solution that contains 0.06 M Na<sub>2</sub>HPO<sub>4</sub> and 0.12 M KH<sub>2</sub>PO<sub>4</sub>, given that pK<sub>a</sub>(2) for phosphoric acid is 7.21.

$$\begin{aligned} \text{pH} &= \text{pK}_a + \log \frac{[\text{A}^-]}{[\text{HA}]} \\ &= 7.21 + \log (0.06/0.12) = 6.91 \end{aligned}$$

**Check: if more of acid than base, then pH is lower than pK<sub>a</sub>!**

(b) Calculate the pH and pH change when 0.8 g of NaOH is added to 400 mL to the above solution.

*NaOH neutralises the acid according to:*



*Need to calculate new conc's of buffer components.*

*Initial moles:*

$$n(\text{H}_2\text{PO}_4^-) = c \times V = 0.12 \text{ mol L}^{-1} \times (400\text{mL}/1000\text{mL/L}) = 48 \text{ mmol}$$

$$n(\text{HPO}_4^{2-}) = 1/2 \times n(\text{H}_2\text{PO}_4^-) = 24 \text{ mmol}$$

*(Note: 1/2 as much H<sub>2</sub>PO<sub>4</sub><sup>-</sup> as HPO<sub>4</sub><sup>2-</sup>)*

*Change in moles (due to NaOH):*

$$n(\text{NaOH}) = m(\text{NaOH})/M(\text{NaOH}) = 0.8 \text{ g}/40 \text{ g mol}^{-1} = 20 \text{ mmol}$$

*Final moles*

$$n(\text{H}_2\text{PO}_4^-) = 48 - 20 = 28 \text{ mmol}$$

$$n(\text{HPO}_4^{2-}) = 24 + 20 = 44 \text{ mmol}$$

*So final conc's:*

*(Note: 1 mmol = 1/1000 mol so the 1000's cancel)*

$$c(\text{H}_2\text{PO}_4^-) = n/V = 28/1000 \text{ mol} / (400\text{mL}/1000\text{mL/L}) = 0.0700 \text{ mol L}^{-1}$$

$$c(\text{HPO}_4^{2-}) = n/V = 44/1000 \text{ mol} / (400\text{mL}/1000\text{mL/L}) = 0.1100 \text{ mol L}^{-1}$$

*Therefore:*

$$\begin{aligned} \text{pH} &= \text{pK}_a + \log \frac{[\text{A}^-]}{[\text{HA}]} \\ &= 7.21 + \log (0.1100/0.0700) = 7.41 \end{aligned}$$

**Check: if more base than acid, then pH is higher than pK<sub>a</sub>!**

*And |pH change| = 7.41 – 6.91 = 0.50 pH units*

(c) Briefly explain why the pH change is so small.

*The added base reacts with H<sub>2</sub>PO<sub>4</sub><sup>-</sup> (the acid in the buffer) and, therefore, the concentration of H<sup>+</sup>(aq) is almost unchanged.*

### Question 3.

(a) Fill in the missing entries in the table below for an aqueous solution at 25 °C.

$[H^+]$ /mol L <sup>-1</sup>	pH	$[OH^-]$ /mol L <sup>-1</sup>	pOH
$= 10^{-pH}$ $= 3.5 \times 10^{-3}$	$= 14 - pOH$ $= 2.5$	$= 10^{-pOH}$ $= 3.2 \times 10^{-12}$	11.5

or  $= K_w/[H^+] = 10^{-14}/3.5 \times 10^{-3}$

$= 2.9 \times 10^{-12}$  (so precise answers will depend on fig's carried & route taken)

(b) The following questions refer to the titration curve shown below right.

(i) Estimate the  $pK_a$  of the substance being titrated.

$pK_a = pH @ \frac{1}{2} \text{ equiv. vol} \sim 8.7$   
(any number a bit less than 9 would do)

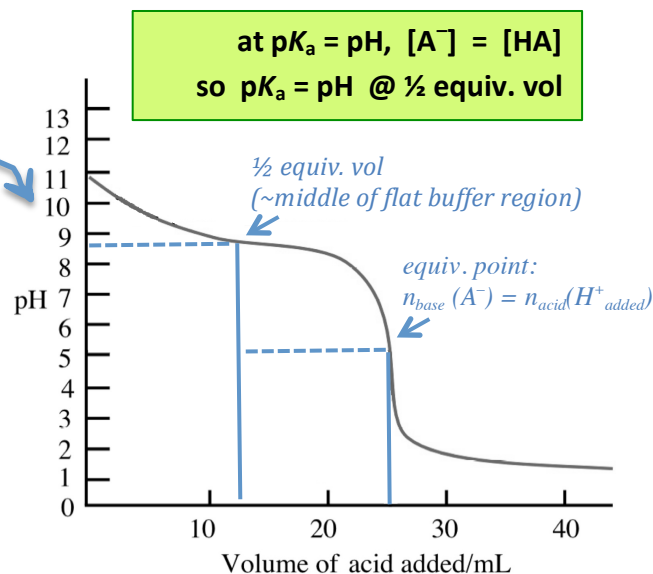
(ii) What class of titration does this curve typify?  
Circle your answer in the list below.

strong base/strong acid

strong base/weak acid

**weak base/strong acid**

weak base/weak acid



(c)  $pK_w = 11.0$  for superheated  $H_2O$  at 250 °C. Calculate the pH of neutral water at this temperature.



**At neutral pH:  $[H^+] = [OH^-]$**

In neutral  $H_2O$ :  $[H^+] = [OH^-]$

so:  $K_w = [H^+][OH^-] = [H^+]^2$

and  $pK_w = -\log([H^+]^2) = -2 \log[H^+] = 2 \times pH$

Hence:  $pH = \frac{1}{2} pK_w = \frac{1}{2} \times 11.0 = 5.5$  for neutral water at 250 °C