

You must show your work for full credit.

### Exp 3 and 4 Questions

1. A  $\text{NH}_3$  solution is being titrated by HCl. (2 pt)

a) Write the reaction that can be used to determine the pH at the equivalence point.



b) Is the solution at the equivalence point acidic, neutral or basic?

**acidic** – above reaction produces  $\text{H}^+$

2. If an unknown weak acid is being titrated by a strong base and a sharp increase in pH occurs at 24.1 mL. A student suspects this corresponds to a second equivalence point for a diprotic acid. Where would he expect the first equivalence point (whether it can be observed or not) to be? (1 pt)

The first equivalence point should occur at half of the second equivalence point. If the acid is diprotic, the first equivalence point would be at **12.05 mL**.

### Additional Titration problems

3. An acetic acid solution (25.0 mL of 0.0750 M) is being titrated by 0.0500 M NaOH.

a) What is the volume of NaOH needed to reach equivalence point in this titration? (1 pt)

$$n(\text{HC}_2\text{H}_3\text{O}_2) = n(\text{NaOH}) \text{ or } [\text{HC}_2\text{H}_3\text{O}_2]V(\text{HC}_2\text{H}_3\text{O}_2) = [\text{NaOH}]V(\text{NaOH})$$

$$\text{or } V(\text{NaOH}) = (25.0)(0.0750 \text{ M}) / (0.0500 \text{ M}) = \mathbf{37.5 \text{ mL}}$$

b) Calculate the pH (to 3 sig fig) after adding 20.0 mL of NaOH. ( $K_a = 1.76 \times 10^{-5}$ ) (3 pts)

Because 20.0 mL < 37.5 mL, this is before the equivalence point in the “buffer” region. We need to determine the moles of and through a mole table and use the H-H equation to determine the pH.

Reaction	$\text{HC}_2\text{H}_3\text{O}_2(\text{aq})$	$+ \text{OH}^-(\text{aq})$	$\leftrightarrow$	$\text{C}_2\text{H}_3\text{O}_2^-(\text{aq})$	$+ \text{H}_2\text{O}(\text{l})$	$\text{pH} = \text{p}K_a + \log[n(\text{A}^-)/n(\text{HA})]$
initial	0.001875	0.00100		0		$\text{p}K_a = -\log(1.8 \times 10^{-5})$
change	-0.00100	-0.00100		+0.00100		$\text{p}K_a = 4.754$
full right	0.000875	0		0.00100		$\text{pH} = 4.754 + \log[(0.00100)/(0.000875)]$
						$\text{pH} = 4.754 + \log(1.143) = \mathbf{4.81}$

grading: give partial credit if correctly calculate moles, set up mole table or use the H-H equation

(continued on back)

c) Which of these indicators is best for observing the equivalence point? You do not need to calculate the equivalence point pH, but explain your answer (1 pt)

Bromcresol Green ( $pK_a = 4.7$ ) Bromthymol Blue ( $pK_a = 6.7$ ) o-Cresolphthalein ( $pK_a = 9.1$ )

*At the equivalence point, we are left with  $C_2H_3O_2^-$ , which is a weak base. Thus, we expect a  $pH > 7$ .*

### Solubility Problem

4. Calculate the molar solubility of silver chromate in water.  $K_{sp}(Ag_2CrO_4) = 1.12 \times 10^{-12}$  (2 pts)

Reaction:  $Ag_2CrO_4(s) \leftrightarrow 2Ag^+(aq) + CrO_4^{2-}(aq)$  which can be solved by using an ICE table

initial	0	0
change	+2S	+S (where S = molar solubility)
equil.	2S	S

$$K_{sp} = 1.12 \times 10^{-12} = [Ag^+]^2[CrO_4^{2-}] = (2x)^2x \text{ or } x = (1.12 \times 10^{-12}/4)^{1/3} = 6.54 \times 10^{-5} \text{ M}$$

*can give partial credit for setting up ICE table or for using the correct equilibrium equation ( $K_{sp}$  equation).*