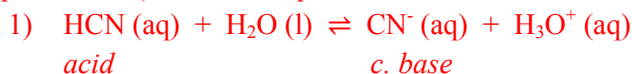


## CHM152 – Chapter 15: Applications of Aqueous Equilibria – Homework

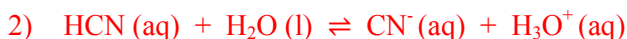
1. What is the pH of a solution prepared by mixing 50.00 mL of 0.10M HCN with 50.00 mL of 0.050 M NaCN? Assume that the volumes are additive and that  $K_a = 4.9 \times 10^{-10}$  for HCN.

Two approaches. 1) Use H-H Equation for buffer solution OR 2) Use ICE table



$$\text{pH} = \text{p}K_a + \log([\text{base}]/[\text{acid}])$$

$$\text{pH} = \text{p}K_a + \log(0.025/0.050) = \mathbf{9.01}$$



$$0.050\text{M}$$

$$0.025$$

$$-x$$

$$+x$$

$$+x$$

$$0.050 - x$$

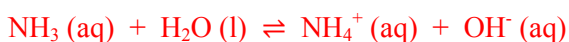
$$0.025 + x$$

$$x$$

$$4.9 \times 10^{-10} = \frac{x(0.025 + x)}{(0.050 - x)} \quad \text{Assume } x \text{ is small} = 4.9 \times 10^{-10} = \frac{x(0.025)}{(0.050)} \quad x = 9.8 \times 10^{-10}$$

$$\text{pH} = -\log(9.8 \times 10^{-10}) = \mathbf{9.01}$$

2. What is the pH of a solution prepared by mixing 50.00 mL of 0.10 M ammonia with 25.00 mL of 0.10 M ammonium chloride? Assume that the volume of the solutions are additive and that  $K_b = 1.8 \times 10^{-5}$  for ammonia.



*base*

*c. acid*

$$0.05000\text{L}(0.10 \text{ M NH}_3)/0.07500\text{L} = 0.067 \text{ M NH}_3$$

$$0.02500\text{L}(0.10 \text{ M NH}_4^+)/0.07500\text{L} = 0.033 \text{ M NH}_4^+$$

$$\text{pH} = -\log(K_w/K_b) + \log([\text{base}]/[\text{acid}])$$

$$\text{pH} = 9.26 + \log(0.067/0.033) = \mathbf{9.56}$$

3. Which one of the following sets of compounds is not characteristic of a buffer?

- $\text{HNO}_3$  with  $\text{NaNO}_3$  Buffers are weak acids with their *c. base* OR weak bases with their *c. acids*
- $\text{NaH}_2\text{PO}_4$  with  $\text{Na}_2\text{HPO}_4$
- $\text{NH}_3$  with  $\text{NH}_4\text{Cl}$
- $\text{CH}_3\text{COOH}$  with  $\text{NaCH}_3\text{COO}$

4. What is the pH of a buffer system made by dissolving 17.42 g of  $\text{KH}_2\text{PO}_4$  and 20.41 g of  $\text{K}_2\text{HPO}_4$  in water to give a volume of 200.0 mL? The  $K_{a2}$  for dihydrogen phosphate is  $6.2 \times 10^{-8}$  and the equilibrium reaction of interest is

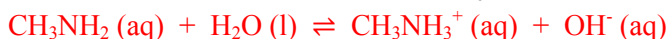


$$17.42 \text{ g KH}_2\text{PO}_4 (1 \text{ mol}/136.09 \text{ g}) = 0.1280 \text{ mol} / 0.2000\text{L} = 0.6400\text{M H}_2\text{PO}_4^- \text{ (acid)}$$

$$20.41 \text{ g K}_2\text{HPO}_4 (1 \text{ mol}/174.18 \text{ g}) = 0.1172 / 0.2000 \text{ L} = 0.5859 \text{ M HPO}_4^{2-} \text{ (c. base)}$$

$$\text{pH} = -\log(6.2 \times 10^{-8}) + \log(0.5859/0.6400) = \mathbf{7.17}$$

5. What is the pH of a buffer system made by dissolving 10.70 g of  $\text{CH}_3\text{NH}_3\text{Cl}$  and 20.00 mL of 12.0 M  $\text{CH}_3\text{NH}_2$  in enough water to make 500.0 mL of solution?  $K_b = 3.7 \times 10^{-4}$  for  $\text{CH}_3\text{NH}_2$ .



$$\text{pH} = 10.57 + \log(0.48/0.31690) = \mathbf{10.75}$$

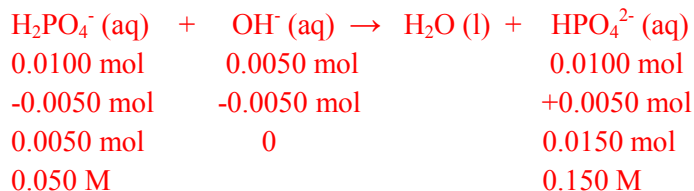
6. What is the  $[\text{CH}_3\text{COO}^-] / [\text{CH}_3\text{COOH}]$  ratio necessary to make a buffer solution with a pH of 4.44?

$$K_a = 1.8 \times 10^{-5} \text{ for CH}_3\text{COOH.}$$

$$4.44 = 4.74 + \log([\text{CH}_3\text{COO}^-]/[\text{CH}_3\text{COOH}]) = -0.30 = \log[\text{CH}_3\text{COO}^-]/[\text{CH}_3\text{COOH}] \quad 10^{-0.30} = \mathbf{0.50:1 \text{ ratio}}$$

7. What is the resulting pH when 0.00500 moles of KOH is added to 0.100 L of a buffer solution that is 0.100 M in  $\text{H}_2\text{PO}_4^-$  and 0.100 M  $\text{HPO}_4^{2-}$  and the  $K_{a2} = 6.2 \times 10^{-8}$ ?

Since a strong base is added, the acid part of the buffer solution

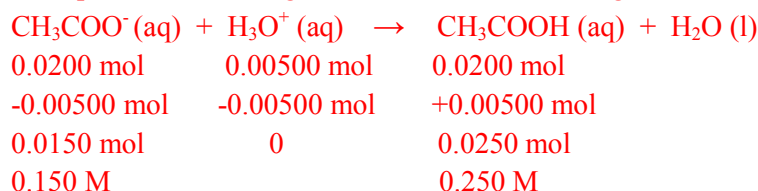


$$\text{pH} = 7.21 + \log(0.15/0.050) = 7.69$$

8. What is the **change** in pH when 0.00500 moles of HCl is added to 0.100 L of a buffer solution that is 0.200 M in  $\text{CH}_3\text{COOH}$  and 0.200 M  $\text{NaCH}_3\text{COO}$ ? The  $K_a$  for acetic acid is  $1.8 \times 10^{-5}$ .

Find the initial pH of the buffer before the acid is added: Since the weak acid and its c. base are equal molarities and volumes, the  $\text{pH} = \text{p}K_a = 4.74$ .

Now find the pH after the strong acid is added. Since strong acid is being added, the c. base reacts with it.



$$\text{pH} = 4.74 + \log(0.150/0.250) = 4.52 \quad \text{change} = \text{pH went down } 0.22$$

9. Describe whether the *resulting* solution when the two reagents are mixed will be acidic, basic or neutral at the equivalence point.
- $\text{CH}_3\text{COOH}$  titrated with NaOH **basic**
  - HF titrated with KOH **basic**
  - HCl titrated with NaOH **neutral**
  - $\text{CH}_3\text{NH}_2$  titrated with HCl **acidic**

10. Formic acid ( $\text{HCOOH}$ ,  $K_a = 1.8 \times 10^{-4}$ ) is the principal component in the venom of stinging ants. What is the molarity of a formic acid solution if 25.00 mL of the formic acid solution requires 29.80 mL of 0.0567 M NaOH to reach the equivalence point?

Use stoichiometry to solve:

$$0.02980 \text{ L} (0.0567 \text{ mol NaOH} / 1 \text{ L}) (1 \text{ mol formic acid} / 1 \text{ mol NaOH}) = 1.69 \times 10^{-3} \text{ mol formic acid} / 0.0250 \text{ L} = 0.0676 \text{ M formic acid}$$

11. What is the pH at the equivalence point of a weak base-strong acid titration if 20.00 mL of NaOCl requires 28.30 mL of 0.50 M HCl?  $K_a = 3.0 \times 10^{-8}$  for HOCl.

$$0.02830 \text{ L} (0.50 \text{ mol HCl} / 1 \text{ L}) (1 \text{ mol OCl}^- / 1 \text{ mol HCl}) / 0.04830 \text{ L} = 0.293 \text{ M OCl}^-$$

Solve the ICE table for  $x = 9.4 \times 10^{-5}$   $\text{pH} = 4.03$

12. Consider the titration of 200.0 mL of 0.150 M hydrofluoric acid in a flask with 0.175 M sodium hydroxide in a burette and answer the following: ( $K_a = 3.5 \times 10^{-4}$  for HF (aq))

- a. How many mL of NaOH must be added to reach the equivalence point?

171 mL

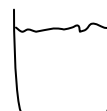
- b. What is the pH after 38.5 mL of NaOH is added?

$$\text{pH} = 2.919 = 2.92$$

- c. What will the resulting solution look like at the equivalence point?

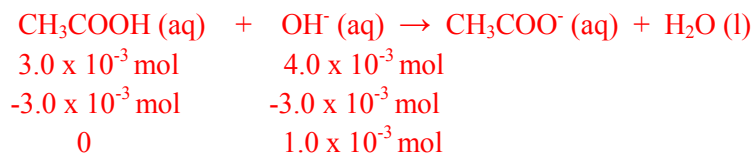
$\text{Na}^+$  and  $\text{F}^-$  are the only ions that should be in solution.

All the  $\text{H}^+$  and  $\text{OH}^-$  are reacted to form water.



13. What is the pH of a solution made by mixing 30.00 mL of 0.10 M acetic acid with 40.00 mL of 0.10 M KOH? Assume that the volumes of the solutions are additive.  $K_a = 1.8 \times 10^{-5}$  for acetic acid.

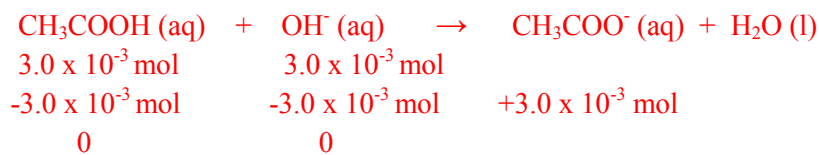
There is more OH<sup>-</sup> than acetic acid so find the moles of OH<sup>-</sup> left and determine pH from that.



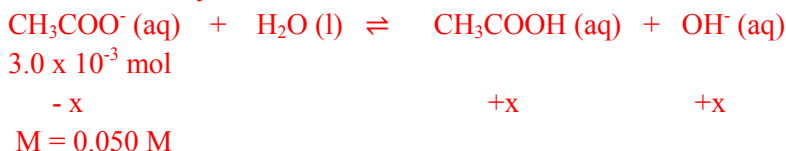
[OH<sup>-</sup>] left = 0.0143 M so pOH = 1.85 and **pH = 12.15**

Acetate ion can further react with water to make acetic acid and hydroxide ion. So more hydroxide ion is produced but the amount is so small it is negligible.

14. What is the pH of a solution made by mixing 30.00 mL of 0.10 M acetic acid with 30.00 mL of 0.10 M KOH? Assume that the volumes of the solutions are additive.  $K_a = 1.8 \times 10^{-5}$  for acetic acid.

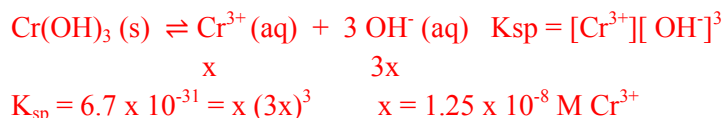


All the acetic acid and hydroxide ion are consumed to form  $3.0 \times 10^{-3}$  mol of acetate ion. It can react with water:

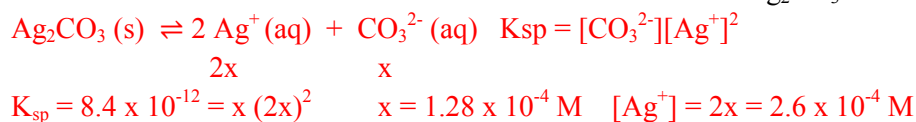


Use  $K_b$ :  $5.6 \times 10^{-10} = x^2 / 0.050$   $x = 5.3 \times 10^{-6}$  M OH<sup>-</sup> **pH = 8.72**

15. What is the chromium ion concentration for a saturated solution of Cr(OH)<sub>3</sub> if the  $K_{sp}$  for Cr(OH)<sub>3</sub> is  $6.7 \times 10^{-31}$ ?



16. What is the silver ion concentration for a saturated solution of Ag<sub>2</sub>CO<sub>3</sub> if the  $K_{sp}$  for Ag<sub>2</sub>CO<sub>3</sub> is  $8.4 \times 10^{-12}$ ?



17. What is the most soluble salt in the following set?

- AgCN with  $K_{sp} = 6.0 \times 10^{-17}$
- Al(OH)<sub>3</sub> with  $K_{sp} = 1.9 \times 10^{-33}$
- Fe(OH)<sub>3</sub> with  $K_{sp} = 2.6 \times 10^{-39}$
- Sn(OH)<sub>2</sub> with  $K_{sp} = 1.6 \times 10^{-19}$

18. Calculate the solubility (in g/L) of silver carbonate in water at 25°C if the  $K_{sp}$  is  $8.4 \times 10^{-12}$ .

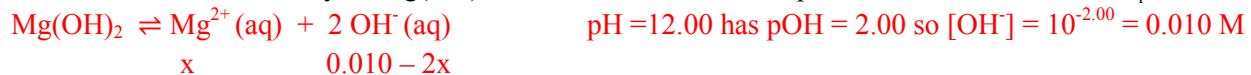
Refer to question #16 for the molar solubility.  $x = 1.28 \times 10^{-4}$  M

For solubility in g/L multiply the molar solubility by the molar mass:  $1.28 \times 10^{-4} \text{ M}(275.78 \text{ g/mol}) = 3.5 \times 10^{-2} \text{ g/L}$

19. What is the molar solubility of CaF<sub>2</sub> in 0.10 M NaF solution at 25°C if the  $K_{sp}$  is  $1.5 \times 10^{-10}$ .

$x = 1.5 \times 10^{-8} \text{ M}$

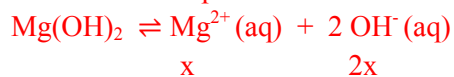
20. What is the molar solubility of  $\text{Mg}(\text{OH})_2$  in a basic solution with a pH of 12.00 at 25°C if the  $K_{\text{sp}}$  is  $5.6 \times 10^{-12}$ .



$$5.6 \times 10^{-12} = x(0.010 - 2x)^2 \quad \text{Assume } x \text{ is small}$$

$$5.6 \times 10^{-12} = x(0.01)^2 \quad x = 5.6 \times 10^{-8} \text{ M}$$

If you wanted to have a little fun, you could calculate the molar solubility of magnesium hydroxide in water and see how a basic pH affects the solubility!



$$5.6 \times 10^{-12} = x(2x)^2 \quad 5.6 \times 10^{-12} = 4x^3 \quad x = 1.1 \times 10^{-4} \text{ M}$$

So having a high pH means more  $\text{OH}^-$  around in the solution which is a common ion. It decreases the solubility of  $\text{Mg}(\text{OH})_2$ !

21. What will occur when calcium nitrate is added to a saturated solution of calcium hydroxide?

Adding a common ion always decreases the solubility of the insoluble compound. In this case calcium hydroxide is slightly soluble in water and by adding calcium nitrate ( $\text{Ca}^{2+}$  is the common ion) the calcium hydroxide becomes even less soluble.