

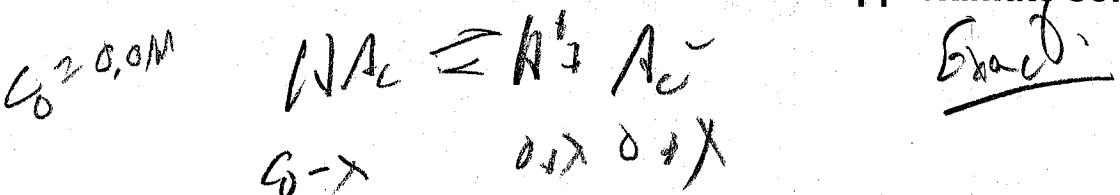
Chap. 8
Mar. 26

Dissociation of Acetic Acid
Slide #11

Example: Acetic Acid - $K_a = 1.8 \times 10^{-5}$

(a) Calculate the pH and percent dissociation
of 0.01 M Acetic Acid - Result for Part (b) is on the slide

We will calculate both the exact and approximate solutions



$$K_a = 1.8 \times 10^{-5} = \frac{x \cdot x}{C_0 - x} \rightarrow x^2 = C_0 K_a = 1.8 \times 10^{-5} \times 0.01 = 1.8 \times 10^{-7}$$
$$x^2 + K_a x - C_0 K_a = 0 \rightarrow [H^+] = \sqrt{1.8 \times 10^{-5} \times 0.01} = 4.24 \times 10^{-4}$$
$$\text{pH} = 3.37$$

$$\text{Approx} \quad K_a = 1.8 \times 10^{-5} = \frac{x \cdot x}{C_0 - x} \approx \frac{x^2}{C_0}$$

$$x^2 = C_0 K_a = (0.01)(1.8 \times 10^{-5}) = 1.8 \times 10^{-7}$$

$$x = \sqrt{1.8 \times 10^{-7}} = 4.24 \times 10^{-4}$$
$$\text{pH} = -\log(4.24 \times 10^{-4}) = 3.37$$

Discuss on next pg

Sik #11

Cont'd.

$$W_{A_0} \geq W_0 A_0$$

C₀

$$x = 4.24 \times 10^{-4} m$$

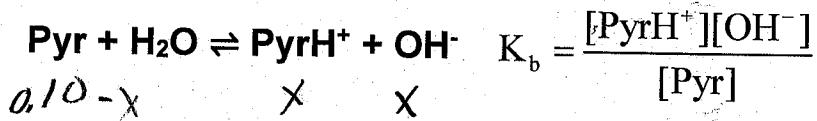
% Des^{50%}

$$\frac{E_A - 1}{\sum E A_0 T_0} \times 100 = \frac{4.24 \times 10^{-4} m}{0.01 m} \times 100$$
$$= 4.24 \times 10^{-2} \times 100 = 42.4\%$$

Protonation of Pyridine

Slide #14

Example: Pyridine - $K_b = 1.8 \times 10^{-9}$



Calculate the pH and the percent of pyridine molecules which are protonated in a 0.10 M aqueous pyridine solution.

$$K_b = 1.8 \times 10^{-9} = \frac{[\text{PyrH}^+][\text{OH}^-]}{[\text{Pyr}]} = \frac{x \cdot x}{0.10-x} \approx \frac{x^2}{0.1}$$

$$x^2 = 1.8 \times 10^{-9} (0.1) = 1.8 \times 10^{-10}$$

$$x = \sqrt{1.8 \times 10^{-10}} = 1.34 \times 10^{-5} = [\text{OH}^-] = [\text{PyrH}^+]$$

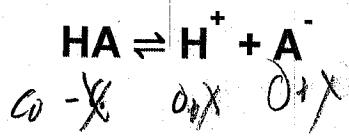
$$\text{pOH} = -\log(1.34 \times 10^{-5}) = 4.13$$

$$\text{pH} = 14 - 4.13 = 9.87$$

$$\% \text{ P}_{\text{at}}: \frac{[\text{PyrH}^+]}{[\text{Pyr}]_0} \times 100 = \frac{1.34 \times 10^{-5}}{0.1} \times 100$$

$$= 0.0134\% \approx 0.013\%$$

Summary: Acid and Base Calculations

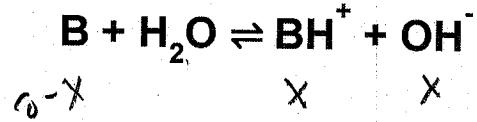


$$K_a = \frac{[H^+][A^-]}{[HA]} = \frac{x \cdot x}{c_0 - x} \approx \frac{x^2}{c_0}$$

$$x = [H^+] = [A^-]$$

$$pH = -\log [H^+] = -\log (x)$$

$$\% \text{ Diss.} = \frac{[x]}{c_0} \times 100$$



$$K_b = \frac{[BH^+][OH^-]}{[B]} = \frac{x \cdot x}{c_0 - x} = \frac{x^2}{c_0}$$

$$pH = -\log (x) = -\log [OH^-]$$

$$pH = 14 - pOH$$

$$\% \text{ Prot.} = \frac{[BH^+]}{c_0} \times 100$$

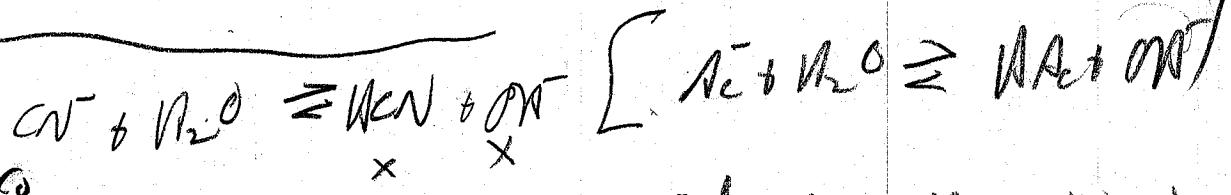
$$x = [BH^+] = [OH^-]$$

Example: $K_a(\text{HCN}) = 4.9 \times 10^{-10}$

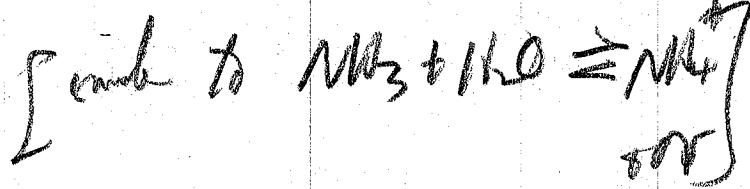
Calculate the pH of a 0.05 M solution of NaCN.



sub to



$$K_b = \frac{[\text{HCN}][\text{OH}^-]}{[\text{CN}^-]}$$



$$= \frac{x \cdot x}{0.05 - x} \approx \frac{x^2}{0}$$

$$2 \times 10^{-5} = \frac{x^2}{0.05} \quad (0.05)$$

$$x^2 = 2 \times 10^{-5} (0.05)$$

$$x^2 = 1 \times 10^{-6}$$

$$x = \sqrt{1 \times 10^{-6}} = 1.0 \times 10^{-3}$$

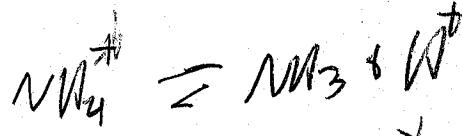
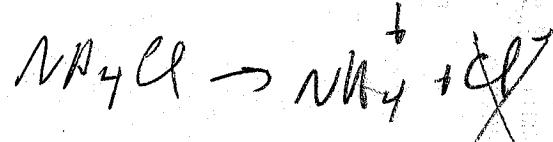
$$\text{pOH} = -\lg(1.0 \times 10^{-3}) = 3.0$$

$$\text{pH} = 14 - 3.0 = 11.0$$

$$\boxed{K_b = \frac{10^{-14}}{K_a} = \frac{10^{-14}}{4.9 \times 10^{-10}}} \\ = 2.0 \times 10^{-5}$$

Example: K_b(NH₃) = 1.8 × 10⁻⁵

Calculate the pH of a 0.20 M solution of NH₄Cl.



$$K_a = \frac{[\text{NH}_3][\text{H}^+]}{[\text{NH}_4^+]} = \frac{x \cdot x}{C_0 - x} = \frac{x^2}{C_0}$$

$$5.6 \times 10^{-10} = \frac{x^2}{0.2}$$

$$x^2 = 1.12 \times 10^{-10}$$

$$x = \sqrt{1.12 \times 10^{-10}} = 1.06 \times 10^{-5}$$

$$\text{pH} = -\log(1.06 \times 10^{-5}) = 4.98 = 5.0$$

$$\boxed{K_a = \frac{1 \times 10^{-10}}{1.8 \times 10^{-5}} = 5.6 \times 10^{-10}}$$

Example: HAc - K_a = 1.8 × 10⁻⁵ $pK_a = -\log(1.8 \times 10^{-5}) = 4.74$

Consider 1 Liter of a solution containing 0.50 M HAc and 0.50 M NaAc. $\rightarrow \text{HAc} + \text{NaAc}$ (0.5M)

(a) Calculate the pH and [H⁺] of the above solution.

(b) Calculate the pH of the above solution after the addition of 0.01 mol of HCl.

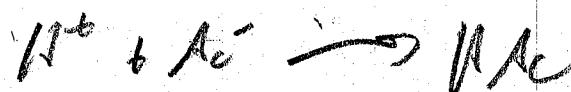
(c) Calculate the pH of the above solution after the addition of 0.01 mol of NaOH.

$$[\text{HAc}] = 0.5 \text{ M} \quad [\text{Ac}^-] = 0.5 \text{ M}$$

$$\text{pH} = pK_a + \log \frac{[\text{Ac}^-]}{[\text{HAc}]} = 4.74 + \log \frac{0.5}{0.5} = 4.74$$

$$[\text{H}^+] = 10^{-4.74} = 1.8 \times 10^{-5}$$

(b) Init: $\text{HAc} = 0.5 \text{ M} / 2 \times 1/2 = 0.5 \text{ M}$
 $\text{NaAc} = 0.5 \text{ M} / 2 \times 1/2 = 0.5 \text{ M}$



init	0.01	0.50	0.50
chg	-0.01	-0.01	+0.01
fin	~0	0.49	0.51

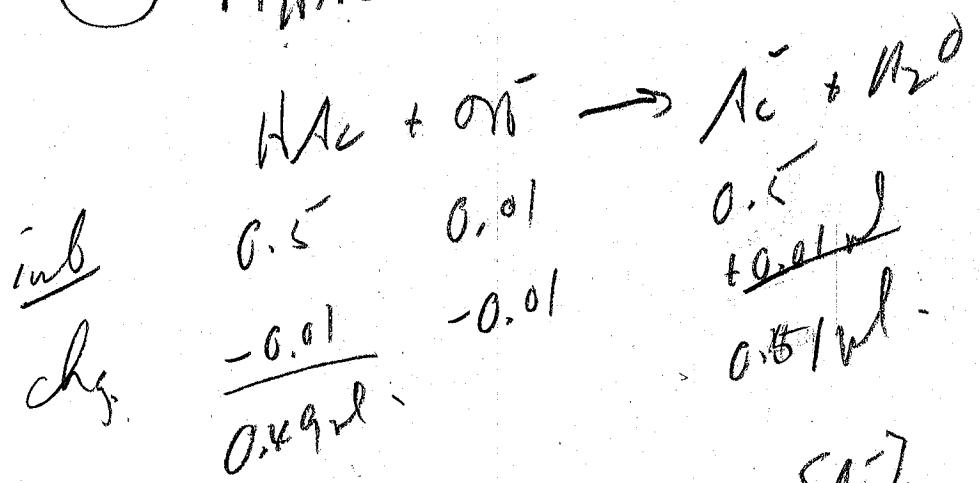
$$[\text{Ac}^-] = 0.49 \quad [\text{HAc}] = 0.51$$

$$\text{pH} = pK_a + \log \frac{[\text{Ac}^-]}{[\text{HAc}]} = 4.74 + \log \frac{0.49}{0.51} = 4.74 - 0.02 = 4.72$$

Stich 35 Sx. (Cbd'l)

(2)

c) $\text{im} \quad n_{\text{NaO}} = 0,5 \text{ mol} \quad n_{\text{Ac}^-} = 0,1$



$$\text{pH} = \text{pK}_a + \log \frac{[\text{Ac}^-]}{[\text{HAc}]} = 4,74 + \log \frac{(0,5)}{0,49}$$

$$= 4,74 + 0,02 = 4,76$$

NH₄⁺/NH₃ Buffer Example

Slide #40

Example: NH₃ - K_b = 1.8 × 10⁻⁵

$$PK_b = -\log(1.8 \times 10^{-5}) = 4.75$$

$$pK_a = 14 - 4.75 = 9.25$$

(A) Calculate the pH of a solution prepared by adding 100 mL of 2.0 M NaOH to 500 mL of 1.2 M NH₄Br

(B) Calculate the pH of a solution prepared by adding 100 mL of 2.0 M HCl to 500 mL of 1.2 M NH₃.

(C) Calculate the ratio, [NH₃] / [NH₄⁺], of the above solution if the pH is adjusted to 9.85

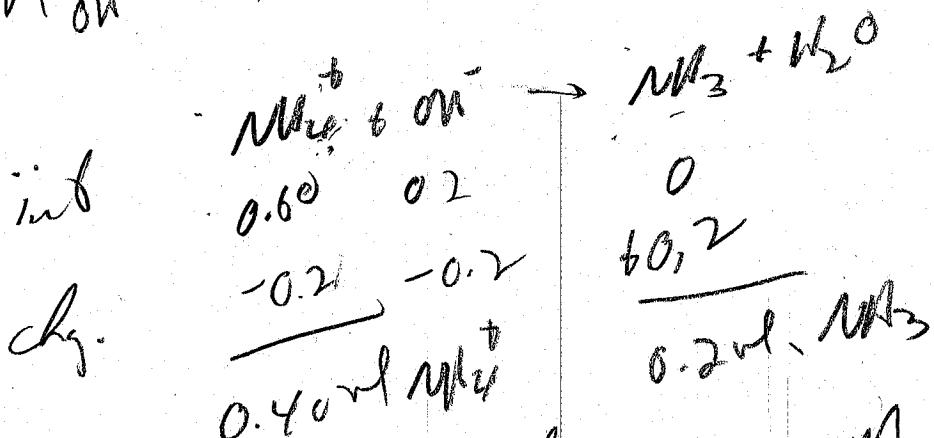
Initial (A)

$$n_{NH_4^+} = 1.2 \text{ M/L} \times 0.50 \text{ L} = 0.60 \text{ mol}$$

$$n_{NH_3} = 0$$

$$n_{OH^-} = 2.0 \text{ M/L} \times 0.10 \text{ L} = 0.20 \text{ mol}$$

$$n_{H^+} = 0$$



$$\frac{\sum NH_4^+ = 0.40 \text{ mol}}{0.62} = 0.67 \text{ M}$$

$$\frac{\sum NH_3 = 0.2 \text{ mol}}{0.62} = 0.33$$

$$pH = pK_a + \log \frac{\sum NH_3}{\sum NH_4^+} = 9.25 + \log \frac{0.33}{0.67}$$

$$= 9.25 + \log(0.5) = 9.25 - 0.30 = 8.95$$

$$\left(= 9.25 + \log \frac{n_{NH_3}}{n_{NH_4^+}} = 9.25 + \log \frac{0.20}{0.40} \right) = 8.95$$

$$\begin{array}{c} NH_4^+ + OH^- \rightarrow NH_3 + H_2O \\ \cancel{pK_b = -\log(1.8 \times 10^{-5}) = 4.75} \\ \cancel{pK_a = 14 - 4.75 = 9.25} \end{array}$$

$$\begin{array}{l} V_{tot} = 100 + 600 \\ = 600 \text{ mL} \\ = 0.6 \text{ L} \end{array}$$

(3)

Slate 40 Ex (Contd)

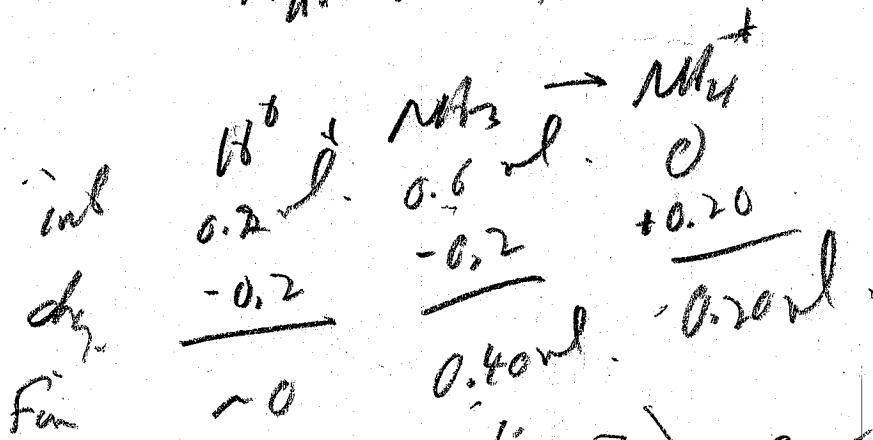
(4)

(B)

$$\text{Init. } N_{NH_3} = 0.2 \text{ mol/l} \times 0.5 \text{ L} = 0.10 \text{ mol}$$

$$N_{NH_3} = 0$$

$$N_{NO} = 2.0 \text{ mol/l} \times 0.1 \text{ L} = 0.20 \text{ mol}$$



$$\begin{aligned}
 \text{pH} &= \text{pK}_a + \log \left(\frac{[NH_3]}{[NH_4^+]} \right) = 9.25 + \log \left(\frac{0.40}{0.20} \right) \\
 &= 9.25 + \log(2) = 9.25 + 0.30 = \underline{\underline{9.55}}
 \end{aligned}$$

Slide 40 (Cont'd.)

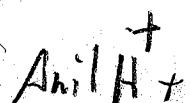
(5)

C) $pH = pK_a + \log \frac{[NH_3]}{[NH_4^+]}$

$$9.85 = 9.25 + \log \frac{[NH_3]}{[NH_4^+]}$$

$$\log \frac{[NH_3]}{[NH_4^+]} = 9.85 - 9.25 = +0.60$$

$$\frac{[NH_3]}{[NH_4^+]} = 10^{0.60} = 3.98 \approx \underline{\underline{4.0}}$$



Review Example: Aniline [Anil] is a weak base with $K_b = 4.3 \times 10^{-10}$

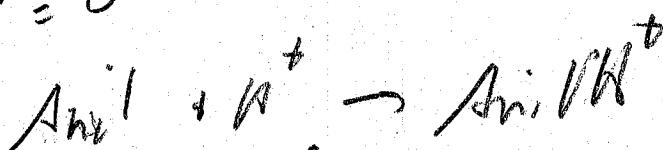
- (A) Calculate the pH of a solution prepared by adding 1.0 L of 3.0 M HCl to 3.0 L of 0.4 M Anil.

- (B) Calculate the pH of a solution prepared by adding 1.0 L of 0.50 M NaOH to 3.0 L of 0.40 M Anilinium Bromide [AnilHBr].

A $\text{Anil}^- = 0.4 \text{ mol/L} \times 3 \text{ L} = 1.2 \text{ mol}$

$$N_{\text{H}^+} = 1 \text{ L} \times 3 \text{ mol} = 0.3 \text{ mol}$$

$$N_{\text{Anil}^+} = 0$$



initial	1.2 mol.	0.3 mol.	0
chg.	-0.3	-0.3	+0.3
	0.9 mol.	0	0.3 mol.

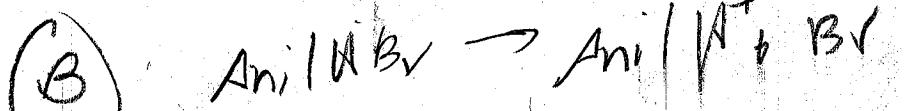
$$[\text{Anil}^-] = \frac{0.9 \text{ mol}}{3.1 \text{ L}}$$

$$[\text{AnilH}^+] = \frac{0.3 \text{ mol}}{3.1 \text{ L}}$$

$$\text{pH} = \text{pK}_a + \log \left(\frac{[\text{Anil}^-]}{[\text{AnilH}^+]} \right) = 4.63 + \log \left(\frac{0.90}{0.3} \right) = 4.63 + 0.48 = 5.11$$

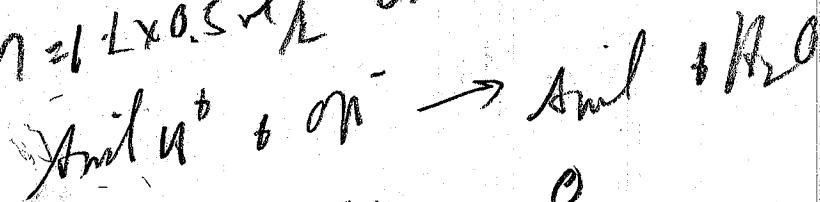
$$\begin{aligned} \text{pK}_b &= -\log(4.3 \times 10^{-10}) \\ &= 9.37 \\ \text{pK}_a &= 14 - 9.37 \\ &= 4.63 \end{aligned}$$

Reaktion (Anilin)



int 0.40 mol/l $\times 3L$
 $= 1.2 \text{ mol}$ 1.2 mol

over n = 1.2 $\times 0.5 \text{ L} = 0.5 \text{ L}$



int	1.2	0.5	0
diff.	<u>-0.5</u>	<u>-0.5</u>	<u>0.5</u>
	0.7 mol		0.5 mol
	anil H^+		anil

$$\begin{aligned}\text{pH} &= \text{p.Ka} + \log \frac{[\text{Stab}]}{[\text{Stoffs}]} = 4.63 + \log \left(\frac{0.5}{0.7} \right) \\ &= 4.63 + (-0.15) \\ &= \boxed{4.48}\end{aligned}$$

Chap. 8
April 4

REVIEW: Homework 8.7

$$pK_a = -\log(1.8 \times 10^{-4}) = 3.74$$

The acid dissociation constant of formic acid [HCOOH] is $K_a = 1.8 \times 10^{-4}$.

A solution is prepared with an initial concentration [HCOOH] = 0.5 M.
Sufficient KOH is added to the solution to raise the pH to 4.25

What are the concentrations, [HCOOH] & [HCOO⁻] of the solution at this pH?

$$\text{initial } [\text{HCOOH}] = 0.50 \text{ M} = [\text{HCOO}^-] + [\text{HCOO}^-]$$

$$\overline{\text{pH} = pK_a + \log \frac{[\text{HCOO}^-]}{[\text{HCOOH}]}}$$

$$4.25 = 3.74 + \log \frac{[\text{HCOO}^-]}{[\text{HCOOH}]}$$

$$\log \frac{[\text{HCOO}^-]}{[\text{HCOOH}]} = 4.25 - 3.74 = +0.51$$

$$\frac{[\text{HCOO}^-]}{[\text{HCOOH}]} = 10^{0.51} = 3.24$$

$$[\text{HCOO}^-] = 3.24 [\text{HCOOH}]$$

$$\overline{[\text{HCOO}^-] + 3.24 [\text{HCOOH}] = 0.5}$$

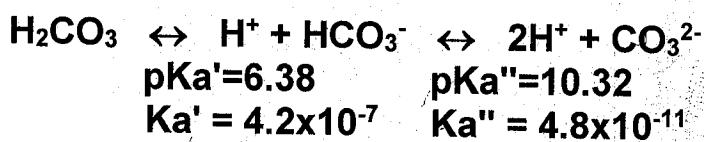
$$4.24 [\text{HCOO}^-] = 0.5$$

$$[\text{HCOO}^-] = \frac{0.5}{4.24} = 0.12 \text{ M}$$

$$\begin{aligned} [\text{HCOO}^-] &= 0.12 - 0.12 \\ &= 0.38 \text{ M} \end{aligned}$$

Chap. 8
April 4

pH of H_2CO_3 and CO_3^{2-}
Slide #50

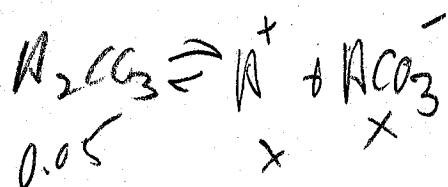


$$K_b'' = \frac{10^{-14}}{K_a''} = \frac{10^{-14}}{4.8 \times 10^{-11}} = 2.1 \times 10^{-4}$$

(a) What is the pH of 0.05 M H_2CO_3 ?

(b) What is the pH of 0.02 M Na_2CO_3 ?

(a)

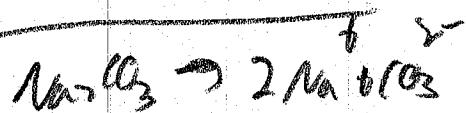


$$\text{K}_a' = 4.2 \times 10^{-7} = \frac{x \cdot x}{0.05}$$

$$x^2 = 2.1 \times 10^{-4}$$

$$x = 1.45 \times 10^{-2}$$

$$\text{pH} = -\log(1.45 \times 10^{-2}) = 3.84$$



(b)



$$K_b'' = \frac{x^2}{0.02} = 2.1 \times 10^{-4}$$

$$x^2 = 4.2 \times 10^{-6}$$

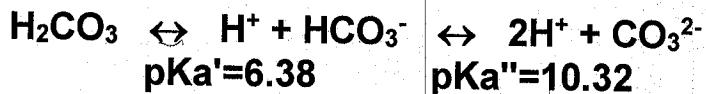
$$x = 2.05 \times 10^{-3}$$

$$= -\log(2.05 \times 10^{-3})$$

$$\text{pOH} = 2.69$$

$$\text{pH} = 14 - 2.69 = 11.31$$

Chap. 8 Carbonic Acid Buffer Questions
 April 4 Slide #54

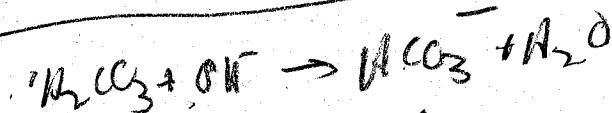


- (1) Calculate the pH of a solution prepared by adding 200 mL of 0.50 M NaOH to 400 mL of 0.40 M H_2CO_2 .
- (2) Calculate the pH of a solution prepared by adding 500 mL of 0.70 M NaOH to 500 mL of 0.40 M H_2CO_2 .
- (3) What ratio of $[\text{H}_2\text{CO}_3]/[\text{HCO}_3^-]$ would be required for a buffer to have $\text{pH} = 5.70$

Q) In. b. $n_{\text{H}_2\text{CO}_3} = 0.4\text{L} \times 0.4\text{M}/2 = 0.16\text{ mol}$,

$$n_{\text{NaOH}} = 0.2\text{L} \times 0.5\text{M}/1 = 0.10\text{ mol}$$

$$\underline{n_{\text{HCO}_3^-} = 0}$$



In. b.	0.16	0.10	$\xrightarrow{+0.10}$
chg.	$\underline{-0.16}$	$\underline{-0.10}$	$\underline{+0.10}$

0.06 mol 0.10 mol $\underline{\text{HCO}_3^-}$



$$\text{pH} = \text{pK}_a' + \log \frac{0.10}{0.06} = 6.38 + 0.22$$

$$= 6.60$$

Carbonic Acid Buff (Cont'd)

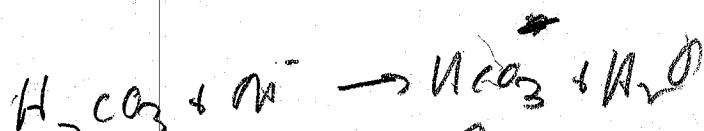
Q.

Int $N_{Na_2CO_3} = 0.50 \text{ L} \times 0.10 \text{ M/L} = 0.05 \text{ mol}$

$$N_{Na_2CO_3^-} = 0$$

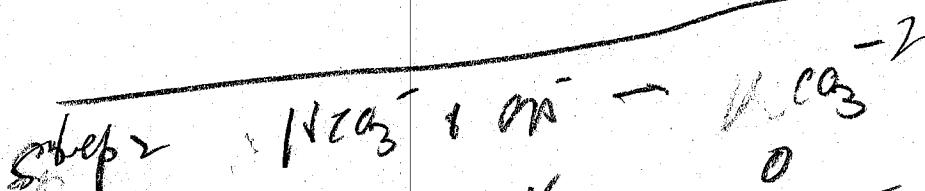
$$N_{NaHCO_3} = 0.50 \text{ L} \times 0.70 \text{ M/L} = 0.35 \text{ mol}$$

Step 1



Int $0.20 \text{ mol} - 0.35 \text{ mol} = 0$

$$\begin{array}{r} -0.2 \\ \hline 0 \end{array} \quad \begin{array}{r} -0.2 \\ \hline 0.15 \end{array} \quad \begin{array}{r} +0.2 \\ \hline 0.2 \text{ mol} \end{array}$$



Int $0.2 \text{ mol} - 0.15 \text{ mol} = 0$

Eq $\begin{array}{r} -0.15 \\ \hline 0 \end{array} \quad \begin{array}{r} +0.15 \\ \hline 0.15 \text{ mol} \end{array}$

$$pH = pK_a'' + \log \frac{[CO_3^{2-}]}{[NaHCO_3^-]}$$

$$= 10.32 + \log \frac{0.15}{0.05}$$

$$= 10.32 + 0.48 = 10.80$$

Carbon Acid Buff (Cont.)

(C)

$$pH = pK_a' + \log \frac{[B(OH)_3^-]}{[BH_2(OH)_2]}$$

$$5.70 = 6.38 + \log \frac{[B(OH)_3^-]}{[BH_2(OH)_2]}$$

$$\log \frac{[B(OH)_3^-]}{[BH_2(OH)_2]} = 5.70 - 6.38 = -0.68$$

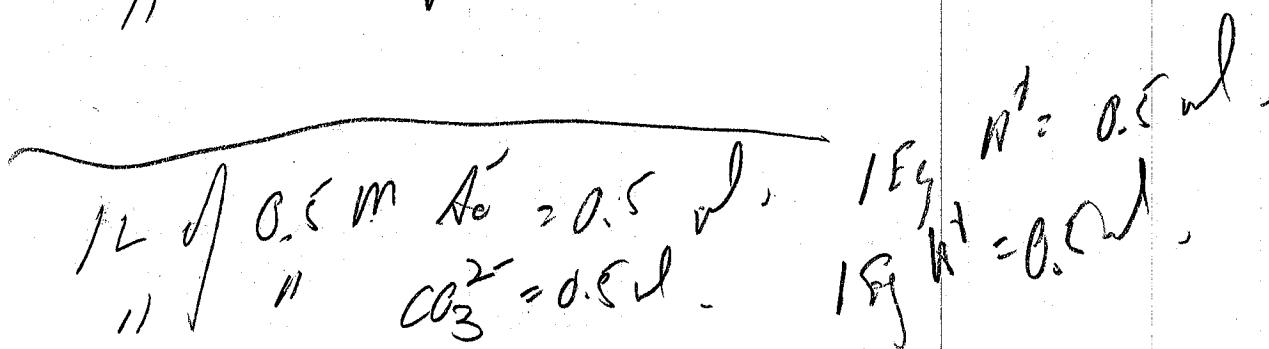
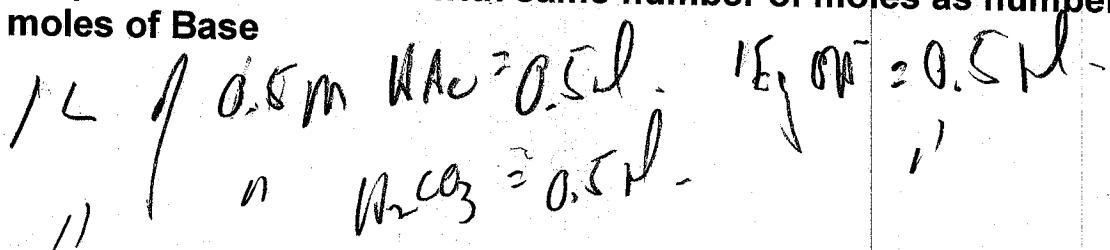
$$\frac{[B(OH)_3^-]}{[BH_2(OH)_2]} = 10^{-0.68} = 0.21$$

$$\frac{[BH_2(OH)_2]}{[B(OH)_3^-]} = \frac{1}{0.21} = 4.8$$

1 Equivalent Definition

Base: 1 equiv. = amount of OH^- with same number of moles as number of moles of Acid

Acid: 1 equiv. = amount of H^+ with same number of moles as number of moles of Base

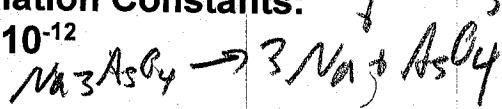


$$pK_a' = \log(16 \times 10^{-3}) = 2.22$$

The following independent questions are on pH calculations in solutions of Arsenic Acid (H_3AsO_4) and its various anions.

H_3AsO_4 is a triprotic acid with Acid Dissociation Constants:

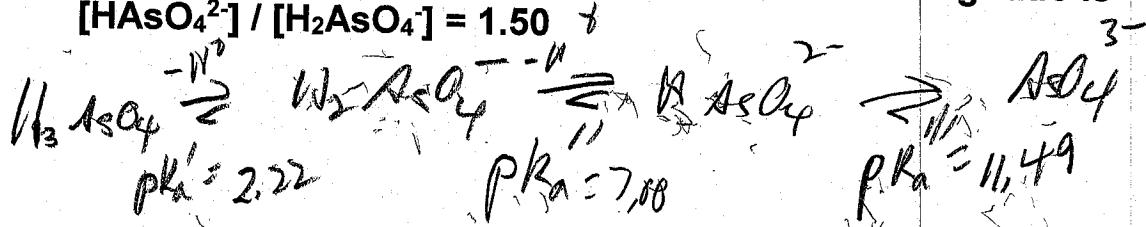
$$K_a' = 6.0 \times 10^{-3}, K_a'' = 1.0 \times 10^{-7}, K_a''' = 3.2 \times 10^{-12}$$



- (C) Calculate the pH of a solution prepared by mixing 3.0 L of 0.40 M Na_3AsO_4 with 1.0 L of 0.80 M HNO_3 .

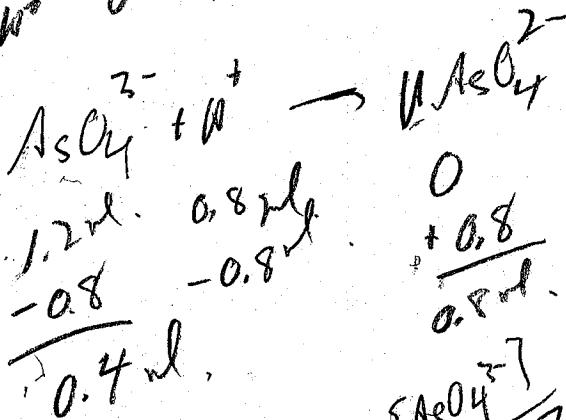
- (D) Calculate the pH of a solution prepared by mixing 3.0 L of 0.40 M Na_3AsO_4 with 2.0 L of 0.80 M HNO_3 .

- (E) What is the pH of a solution in which the following ratio is $[HAsO_4^{2-}] / [H_2AsO_4^-] = 1.50$



(C) $N_{AsO_4} = 0.4 \text{ M} \times 3 \text{ L} = 1.2 \text{ M}$

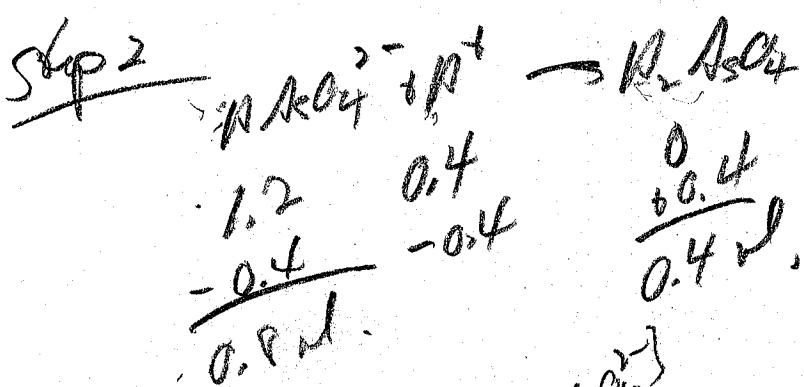
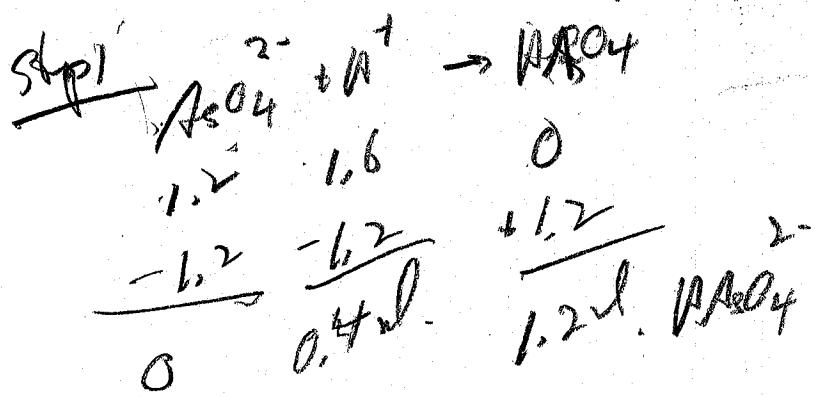
$$N_{H^+} = 0.8 \text{ M} / 2 \times 1 \text{ L} = 0.8 \text{ M}$$



$$pH = pK_a''' + \log \frac{[AsO_4^{3-}]}{[HAsO_4^{2-}]} = 11.49 + \log \frac{0.4}{0.8} = 11.49 - 0.30 = 11.19$$

Ruth PW Solut (Ans)

(A) $\text{Na}_2\text{SO}_4^{2-} = 0.4 \text{ M} / \times 3L = 1.2 \text{ M}$
 $= 1.6 \text{ M}$
 $\eta_{\text{eff}} = 0.8 \times 2$



$$\text{pH} = \text{pK}_a + \log \frac{\text{[HgO}_4^{2-}]_{\text{initial}}}{\text{[HgO}_4^{2-}]}$$

$$= 7.00 + \log \frac{0.8}{0.4}$$

$$= 7.00 + 0.30$$

$$= 7.30$$

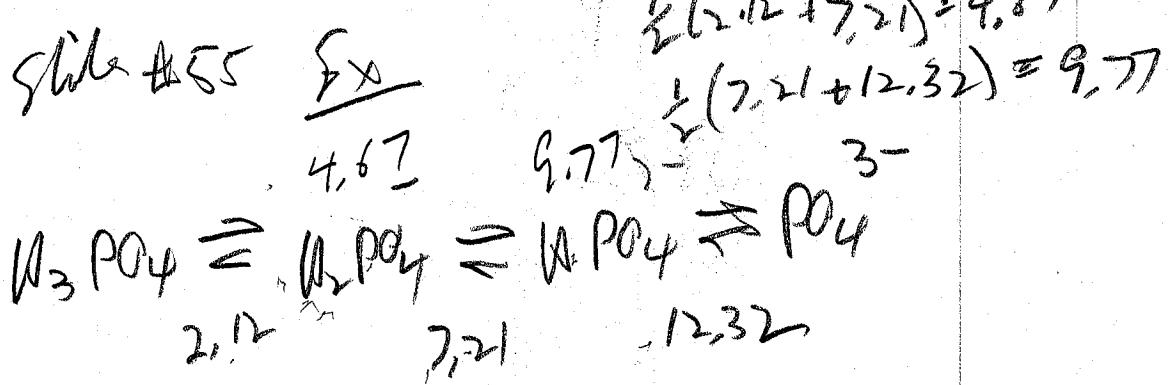
Rev. HW 8.6 (cont.)

(E)

$$\frac{[\text{H}_2\text{AsO}_4^-]}{[\text{H}_2\text{AsO}_4^{\text{H}^+}]} = 1,5 \rightarrow \frac{[\text{H}_2\text{AsO}_4^{\text{H}^+}]}{[\text{H}_2\text{AsO}_4^-]} = \frac{1}{1,5}$$

$$\text{pH} = \text{pK}_a'' + \log \frac{[\text{H}_2\text{AsO}_4^{\text{H}^+}]}{[\text{H}_2\text{AsO}_4^-]}$$

$$= 7,00 + \log(1,5) = 7,00 + 0,18 \\ = 7,18 \approx 7,2$$



a) $pH = 4.67$

b) $pH = 9.77$

c) $pH = pK_a' + \log \frac{[H_2PO_4^-]}{[H_3PO_4]} = 2.12 + \log(1) = 2.12$

d) $pH = pK_a''' + \log \frac{[PO_4^{3-}]}{[H_3PO_4]} = 12.32 + \log 1 = 12.32$

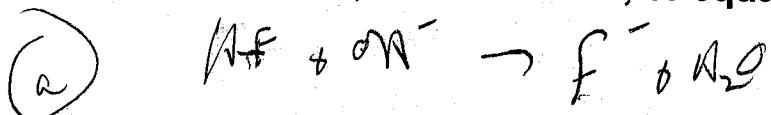
$$pK_a = -\log(7.2 \times 10^{-4}) \\ = 3.14$$

Hydrofluoric Acid, HF, is a weak acid with $K_a = 7.2 \times 10^{-4}$.
One initially has 600 mL of 0.30 M HF,

(a) How many mL of 0.50 M NaOH are required to reach half-way to the equivalence point?

(b) What is the pH of the solution half-way to the equivalence point?

(c) If one adds an additional half equivalence of NaOH, will the pH higher than 7.0, lower than 7.0, or equal to 7.0?



$$N_{\text{HF}} = 0.30 \text{ M} \times 0.6 \text{ L} = 0.18 \text{ mol}$$

$$N_{\text{NaOH}} = \frac{1}{2} N_{\text{HF}} = 0.09 \text{ mol}$$

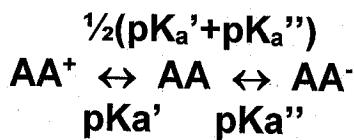
$$C_{\text{NaOH}} = \frac{N_{\text{NaOH}}}{V} \rightarrow V^2 \frac{N_{\text{NaOH}}}{C_{\text{NaOH}}} = \frac{0.09}{0.5 \text{ M}} \\ = 0.18 \text{ L} = 180 \text{ mL}$$

(b) $\text{pH} = pK_a + \log \frac{[\text{F}^-]}{[\text{HF}]}$ =

$$= 3.14 + \log(1) = 3.14$$

(c) $\text{pH} > 7$

pH Dependence of Species Concentrations
Glycine: Slide #70



Example: Glycine - $\text{pK}_a' = 2.34$ $\text{pK}_a'' = 9.60$

(A) What is the pH of 0.10 M neutral Glycine?

(B) If the pH of the above solution is adjusted to 9.0,
what species are present and what are their concentrations?

(C) What are the species present if the pH is adjusted to 4.5?

(A)

$$\text{pH} = 5.97$$



$$\Sigma [\text{Gly}] + [\text{Gly}^-] = 0.10 \text{ M}$$

$$9.00 = \text{pH} = \text{pK}_a + \log \frac{[\text{Gly}^-]}{\Sigma [\text{Gly}]}$$

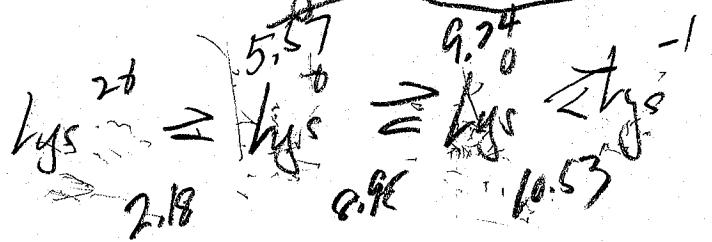
$$\text{for } \frac{[\text{Gly}^-]}{\Sigma [\text{Gly}]} = \frac{9.00 - 9.60}{-0.60} = 0.25$$

$$\frac{[\text{Gly}^-]}{[\text{Gly}]} = 10^{-0.60} = 0.25 \rightarrow [\text{Gly}^-] = 0.25 [\text{Gly}]$$

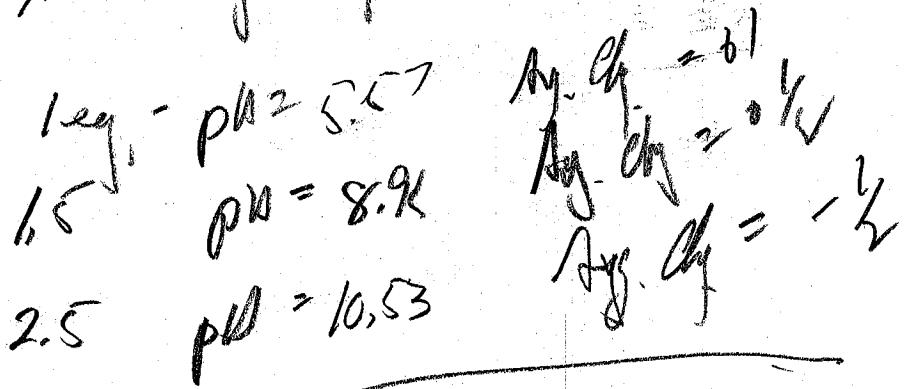
$$\Sigma [\text{Gly}] + 0.25[\text{Gly}] = 0.10 = 1.25 [\text{Gly}]$$

$$[\text{Gly}] = \frac{0.1}{1.25} = 0.08 \text{ M} \quad [\text{Gly}^-] = 0.1 - 0.08 = 0.02$$

Lysine + NH₃ pH 10.74



Total # Egin of OH⁻



pI = 9.74
 2.6g OH⁻ will need to get Ag. Chg = -1
 3