N37 – Acid Base

Weak Problems

Link to YouTube Presentation: https://youtu.be/WwZBnDhd9J0

N37 – Acid Base Weak Problems

Target: I can review Acid Base definitions and perform pH calculations for weak acids/bases

N37 – Acid Base

Yay, ICE Tables are back!

What is the pH of a 0.50 M solution of acetic acid, $HC_2H_3O_2$, $K_a = 1.8 \times 10^{-5}$?

Step #1: Write the dissociation equation

$$HC_2H_3O_2 \leftrightarrows C_2H_3O_2 + H^+$$

What is the pH of a 0.50 M solution of acetic acid, $HC_2H_3O_2$, $K_a = 1.8 \times 10^{-5}$?

Step #2: ICE it!

$$HC_{2}H_{3}O_{2} \leftrightarrows C_{2}H_{3}O_{2}^{-} + H^{+}$$
I 0.50 0 0
C - x +x +x
E 0.50 - x x x

What is the pH of a 0.50 M solution of acetic acid, $HC_2H_3O_2$, $K_a = 1.8 \times 10^{-5}$?

Step #3: Set up the law of mass action

$$HC_2H_3O_2 \leftrightarrows C_2H_3O_2^- + H^+$$

E 0.50 - x x

$$1.8 \times 10^{-5} = \frac{(x)(x)}{(0.50 - x)} \cong \frac{x^2}{(0.50)}$$

Can use the 5% rule because K < 1 and K at least 1000 time smaller than [initial]

What is the pH of a 0.50 M solution of acetic acid, $HC_2H_3O_2$, $K_a = 1.8 \times 10^{-5}$?

Step #4: Solve for x, which is also [H+]

$$HC_2H_3O_2 \leftrightarrows C_2H_3O_2^- + H^+$$

E 0.50 - x x

$$1.8 \times 10^{-5} = \frac{\chi^2}{(0.50)}$$
 [H⁺] = 3.0 x 10⁻³ M

What is the pH of a 0.50 M solution of acetic acid, $HC_2H_3O_2$, $K_a = 1.8 \times 10^{-5}$?

Step #5: Convert [H+] to pH

$$HC_2H_3O_2 \leftrightarrows C_2H_3O_2^- + H^+$$
E 0.50 - x x 3.0 x 10⁻³ M

$$pH = -log(3.0 \times 10^{-3}) = 2.52$$

Reaction of Weak Bases with Water

The base reacts with water, producing its conjugate acid and hydroxide ion:

$$CH_3NH_2 + H_2O \implies CH_3NH_3^+ + OH^- K_b = 4.38 \times 10^{-4}$$

$$K_b = 4.38 \times 10^{-4} = \frac{[CH_3NH_3^+][OH^-]}{[CH_3NH_2]}$$

K_b for Some Common Weak Bases

Many students struggle with identifying weak bases and their conjugate acids. What patterns do you see that may help you?

| Base | Formula | Conjugate Acid | K _b |
|---------------|---|---|-------------------------|
| Ammonia | NH ₃ | NH ₄ + | 1.8 x 10 ⁻⁵ |
| Methylamine | CH ₃ NH ₂ | CH ₃ NH ₃ + | 4.38 x 10 ⁻⁴ |
| Ethylamine | $C_2H_5NH_2$ | C ₂ H ₅ NH ₃ + | 5.6 x 10 ⁻⁴ |
| Diethylamine | $(C_2H_5)_2NH$ | $(C_2H_5)_2NH_2^+$ | 1.3 x 10 ⁻³ |
| Triethylamine | $(C_2H_5)_3N$ | (C ₂ H ₅) ₃ NH ⁺ | 4.0 x 10 ⁻⁴ |
| Hydroxylamine | HONH ₂ | HONH ₃ + | 1.1 x 10 ⁻⁸ |
| Hydrazine | H ₂ NNH ₂ | H ₂ NNH ₃ + | 3.0 x 10 ⁻⁶ |
| Aniline | C ₆ H ₅ NH ₂ | C ₆ H ₅ NH ₃ + | 3.8 x 10 ⁻¹⁰ |
| Pyridine | C_5H_5N | C ₅ H ₅ NH ⁺ | 1.7 x 10 ⁻⁹ |

Reaction of Weak Bases with Water

The generic reaction for a base reacting with water, producing its conjugate acid and hydroxide ion:

$$B + H_2O \leftrightarrows BH^+ + OH^-$$

$$K_b = \frac{[BH^+][OH^-]}{[B]}$$

Yes, all weak bases do this – DO NOT make this more complicated then it needs to be.

What is the pH of a 0.50 M solution of ammonia, NH₃, $K_b = 1.8 \times 10^{-5}$?

Step #1: Write the equation for the reaction

$$NH_3 + H_2O \Rightarrow NH_4^+ + OH^-$$

What is the pH of a 0.50 M solution of ammonia, NH₃, $K_b = 1.8 \times 10^{-5}$?

Step #2: ICE it!

$$NH_3 + H_2O \leftrightarrows NH_4^+ + OH^-$$
I 0.50 0 0
C - x +x +x
E 0.50 - x x x

What is the pH of a 0.50 M solution of ammonia, NH₃, $K_b = 1.8 \times 10^{-5}$?

Step #3: Set up the law of mass action

$$NH_3 + H_2O \leftrightarrows NH_4^+ + OH^-$$
E 0.50 - x x x

$$1.8 \times 10^{-5} = \frac{(x)(x)}{(0.50 - x)} \cong \frac{x^2}{(0.50)}$$

Can use the 5% rule because K < 1 and K at least 1000 time smaller than [initial]

What is the pH of a 0.50 M solution of ammonia, NH₃, $K_b = 1.8 \times 10^{-5}$?

Step #4: Solve for x, which is also [OH-]

$$NH_3 + H_2O \leftrightarrows NH_4^+ + OH^-$$

E 0.50 - x x x

$$1.8 \times 10^{-5} = \frac{\chi^2}{(0.50)}$$
 [OH-] = 3.0 x 10-3 M

What is the pH of a 0.50 M solution of ammonia, NH₃, $K_b = 1.8 \times 10^{-5}$?

Step #5: Convert [OH-] to pH

$$pOH = -log(3.0 \times 10^{-3}) = 2.52$$

 $pH = 14 - pOH = 14 - 2.52 = 11.48$

Remember! Everyone always forgets!

You can convert back and forth from Ka to Kb and vice versa. If you are given Ka for an acid but are doing problems with the acid's conjugate base you can use that Ka to find the Kb that you need.

$$Ka \times Kb = Kw$$

$$Ka \times Kb = (1 \times 10^{-14})$$

Weird Fact!

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

The % ionization for a weak acid INCREASES as concentration DECREASES!

Let's pretend we cut the [] in half

$$Q = \frac{\left|\frac{1}{2}\right|\left|\frac{1}{2}\right|}{\left|\frac{1}{2}\right|} = \frac{1}{2}$$
 (of whatever the original was)

If Q decreases...

Reaction shifts to make more products...

More ionization! Higher % ionization!

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