<u>N-48</u> Weak Acid/Base Equilibria

Weak Acid/Base Equilibria

N - 48

<u>Target</u>

I can use equilibrium expressions and ICE tables to calculate the concentrations of reactants and products at equilibrium for acids and bases, and can use that information to find the pH or other related values.

Link to YouTube Presentation: https://youtu.be/fkc4USA2518





What do chemists mean by <u>WEAK?</u> They do not completely ionize in water. Only a <u>LITTLE BIT</u> will be dissociated.



Dissociation is a reversible reaction right?

So...

We can use equilibrium constants, expressions, ice tables to determine []'s which let us find...

<u>pH values!</u>



- Remember that Keq is just generic. Ka and Kb Could be Kc, Kp, Ksp if you are trying to be specific. So for acid bases use:
 - K_a (for acids)
 - K_b (for bases)

Still $\frac{Products}{Reactants}$ which will be

[Dissociated Ions]

[Undissociated Molecule]

 $\begin{array}{l} HA \leftrightarrow H^{+} + A^{-} \\ BOH \leftrightarrow B^{+} + OH^{-} \\ B + H_{2}O \leftrightarrow BH^{+} + OH^{-} \end{array}$

Practice Problem

Identify Acid/Conj Base/Base/Conj Acid for $HC_2H_3O_2$ (abbreviated as HOAc). Then write the Ka Expression

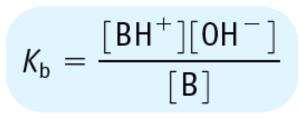
$HC_{2}H_{3}O_{2(aq)} + H_{2}O_{(l)} \leftrightarrow H_{3}O^{+}_{(aq)} + C_{2}H_{3}O_{2}^{-}_{(aq)}$ Acid Base Conj. Acid Conj. base $Ka = \frac{[H_{3}O^{+}][OAc^{-}]}{[HOAc]}$

Size of Ka for Weak Acids

$Ka = \frac{[H_3O^+][OAc^-]}{[HOAc]} = 1.8 \times 10^{-5}$ Why is the Ka so small for acetic acid???

- Small Ka means equilibrium lies to the LEFT
- Reactant Favored not much dissociated
- It is a WEAK acid!
- Weak acids have Ka < 1
 - Leads to low $[H^+] \rightarrow pH$ from 2 6.9





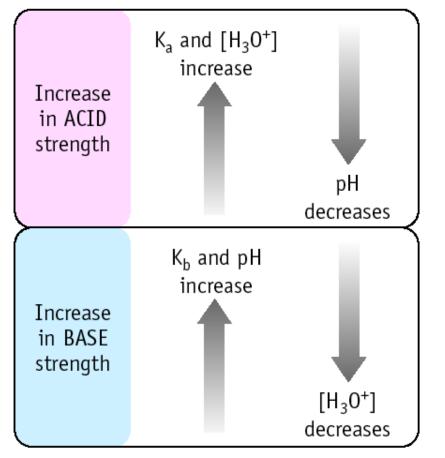
Size of Kb for Weak Bases

$$B(aq) + H_2O(\ell) \implies BH^+(aq) + OH^-(aq)$$

Weak Bases have small Kb values

- Small Kb means equilibrium lies to the LEFT
- Reactant Favored not much dissociated
- It is a WEAK base!
- Weak bases have Kb < 1
 - Leads to low $[OH^-] \rightarrow pH$ from 12 7.1

<u>Relationship between Ka, [H+], pH</u>



Glue Into Your Notes!



You have 1.00 M HOAc. Calculate the equilibrium concentrations of HOAc, H_3O^+ , OAc⁻, and the pH if the Ka = 1.8 x 10⁻⁵.

$HOAc + H_2O \leftrightarrow H_3O^+ + OAc^-$

Step 1: Create an ICE Table
Step 2: Write Ka expression (or Kb expression depending on Q)
Step 3: Solve for x using quadratic or 5% rule if valid.
Step 4: Solve for pH (or pOH depending on the Q)

You have 1.00 M HOAc. Calculate the equilibrium concentrations of HOAc, H_3O^+ , OAc⁻, and the pH if the Ka = 1.8 x 10⁻⁵. HOAc + $H_2O \leftrightarrow H_3O^+ + OAc^-$

Reaction	[HOAc]	[H ₃ O+]	[OAc ⁻]
I	1.00	0.00	0.00
С	-X	+X	+X
E	1.00-x	X	X
5%	1.00	X	X
Answer			

You have 1.00 M HOAc. Calculate the equilibrium concentrations of HOAc, H_3O^+ , OAc⁻, and the pH if the Ka = 1.8 x 10⁻⁵. HOAc + $H_2O \leftrightarrow H_3O^+ + OAc^-$

$$Ka = \frac{[H_3O^+][OAc^-]}{[HOAc]} \qquad 1.8 \ x \ 10^{-5} = \frac{x^2}{1.00}$$

 $x = 4.2 \times 10^{-3}$



You have 1.00 M HOAc. Calc. the equilibrium concentrations of HOAc, H_3O^+ , OAc⁻, and the pH if Ka = 1.8x10⁻⁵. HOAc + $H_2O \leftrightarrow H_3O$ + OAc⁻

Reaction	[HOAc]	[H ₃ O+]	[OAc ⁻]
I	1.00	0.00	0.00
С	-X	+X	+X
E	1.00-x	X	X
5%	1.00	X	x
Answer	1.00	4.2 x 10 ⁻³	4.2 x 10 ⁻³

You have 1.00 M HOAc. Calc. the equilibrium concentrations of HOAc,

 H_3O^+ , OAc^- , and the pH if Ka = 1.8x10⁻⁵. HOAc + $H_2O \leftrightarrow H_3O + OAc^-$

Check that the 5% Rule is Valid!

$$x = \frac{x}{[initial]} \ x \ 100 \ \le 5\%$$

$$x = \frac{4.2 x \, 10^{-3}}{1.00} \, x \, 100 = 0.42\% \quad \checkmark$$

You have 1.00 M HOAc. Calc. the equilibrium concentrations of HOAc, H_3O^+ , OAc⁻, and the pH if Ka = 1.8x10⁻⁵. HOAc + $H_2O \leftrightarrow H_3O + OAc^-$

Now Solve for pH – Don't forget!!!

You have 0.010 M NH₃. Calculate the pH. $K_b = 1.8 \times 10^{-5}$ NH₃ + H₂O \leftrightarrow NH₄⁺ + OH⁻

Reaction	[NH ₃]	[NH ₄ +]	[OH ⁻]
I	0.010	0.00	0.00
С	-X	+X	+X
E	0.010-x	X	X
5%	0.010	X	x
Answer			

You have 0.010 M NH₃. Calculate the pH. $K_b = 1.8 \times 10^{-5}$ NH₃ + H₂O \leftrightarrow NH₄⁺ + OH⁻



$x = 4.2 \times 10^{-4}$

You have 0.010 M NH₃. Calculate the pH. $K_b = 1.8 \times 10^{-5}$ NH₃ + H₂O \leftrightarrow NH₄⁺ + OH⁻

Reaction	[NH ₃]	[NH ₄ +]	[OH ⁻]
I	0.010	0.00	0.00
С	-X	+X	+X
E	0.010-x	x	X
5%	0.010	X	X
Answer	0.010	4.2 x 10 ⁻⁴	4.2 x 10 ⁻⁴

Practice with Weak Acid/Bases You have 0.010 M NH₃. Calculate the pH. $K_h = 1.8 \times 10^{-5}$ $NH_3 + H_2O \leftrightarrow NH_4^+ + OH^-$ Now Calculate pH! $[OH^{-}] = 4.2 \times 10^{-4} M$ $pOH = -\log [OH^{-}] = 3.37$ $pH + pOH = 14 \rightarrow 14 = pH + 3.37$ pH = 10.63



The % dissociation INCREASES as the concentration of the weak acid or weak base DECREASES!





$HA \rightarrow H^+ + A^-$ % dissociation = $\frac{[ions]}{[undissociated]} x 100$ WHICH IS LIKE.... $\mathbf{K}_{\mathbf{a}} = \frac{[H^+][A^-]}{HA}$





HA \rightarrow H⁺ + A⁻ % dissociation = $\frac{[H^+]}{[undissociated]} \times 100$



Let's pretend we take our bottle of acid and double the amount of water.

The concentration of everything is cut in half correct????



So our K_a turns into...

$$\mathbf{Q} = \frac{\left[\frac{1}{2}H^{+}\right]\left[\frac{1}{2}A^{-}\right]}{\left[\frac{1}{2}HA\right]}$$

vs.
$$\frac{[H^+][A^-]}{[HA]}$$
 like it had been.





So... $Q = \frac{\frac{1}{2} [H^+][A^-]}{[HA]}$

So Q = $\frac{1}{2}$ Ka







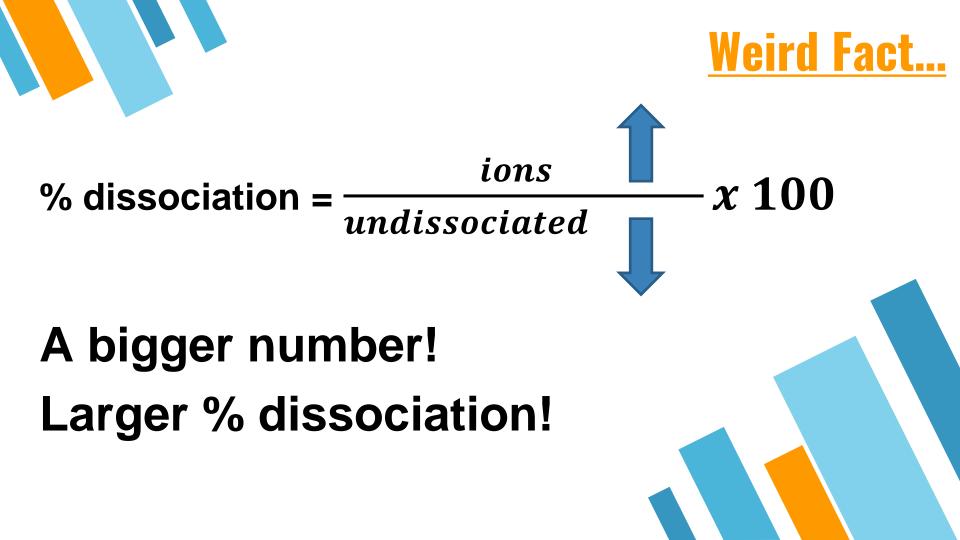
MEANS... Not enough products! SHIFT RIGHT to get to equilibrium! Make more products!

Q < **K**



And the products are... [H⁺] and [A⁻] !

So MORE ions means... HIGHER % dissociation!



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