ACID/BASE THEORY

For the ionization of water : $H_2O + H_2O = H_3O^+ + OH^$ an equilibrium expression can be written: $K_w = [H_3O^+][OH^-] = 1.0 \times 10^{-14}$ (@ 25°C) $- or - K_w = [H^+][OH^-] = 1.0 \times 10^{-14}$ (@ 25°C)

 K_w is know as the **ion product** of water. In **ANY** aqueous solution this equilibrium is always present. One can always obtain the concentration of **H**⁺ from the concentration of **OH**⁻ and vice versa. A strong acid will <u>set</u> the value of [H⁺] and then [OH⁻] can be calculated. A strong base will <u>set</u> the value of [OH⁻] and then [H⁺] can be calculated.

The pH of a solution can be determine by the expression: $pH = -log[H^+]$ which means that $[H^+] = 10^{pH}$

In general, the "p" in pH is really a function: $p(X) = -\log(X)$ but instead of writing p(X) we write pX.

So you can easily calculate values for pOH, pK_w, etc...

This leads to the "-log" form of the above equation: 14 = pH + pOH

The ONLY time that $[H^+] = [OH^-]$ (definition of neutral water) is when they each equal 1.0×10^{-7} M which corresponds to a pH (and pOH for that matter) of 7.00. This is neutral pH and is the pH of pure H₂O (note this is only at 25°C, however). pH values lower than 7 are acidic solutions while pH values higher than 7 are basic.

NOTE: ALL the questions in chapters 6 and 7 will fit into these equilibria and equations compare and contrast the similarities and differences in these two columns

A C I D SB A S E SFor a weak acid (HA) in water:
HA + H2O = H3O^+ + A^-For a weak base (B) in water:
B + H2O = BH^+ + OH^-
$$K_a = \frac{[H^+][A^-]}{[HA]}$$
 $[H^+] = K_a \frac{[HA]}{[A^-]}$ $K_b = \frac{[OH^-][BH^+]}{[B]}$ $[OH^-] = K_b \frac{[B]}{[BH^+]}$ $pH = pK_a + \log \frac{[A^-]}{[HA]}$ $pOH = pK_b + \log \frac{[BH^+]}{[B]}$ $pOH = pK_b + \log \frac{[BH^+]}{[B]}$ For a conjugate acid (BH +) of a
weak base(B) in water:For a conjugate base (A^-) of a
weak acid (HA) in water:BH^+ + H2O = H3O^+ + B
 $K_a = \frac{[H^+][B]}{[BH^+]}$ A^- + H2O = HA + OH^- $K_a = \frac{[H^+][B]}{[BH^+]}$ $[H^+] = K_a \frac{[BH^+]}{[B]}$ $K_b = \frac{[OH^-][HA]}{[A^-]}$ $pH = pK_a + \log \frac{[B]}{[BH^+]}$ $pOH = pK_b + \log \frac{[HA]}{[A^-]}$

NOTICE! You can NOT look up the K_a of BH⁺. You have to calculate it from the K_b of B:

 $K_{\rm a} = \frac{K_{\rm w}}{K_{\rm b}}$

NOTICE! You can NOT look up the K_b of A⁻. You have to calculate it from the K_a of HA:

$$K_{\rm b} = \frac{K_{\rm w}}{K_{\rm a}}$$