

**(14) Today**

Sections 2.7 – 2.11  
Acids and Bases

**Next Class (15)**

Sections 2.7 – 2.11  
Acids and Bases

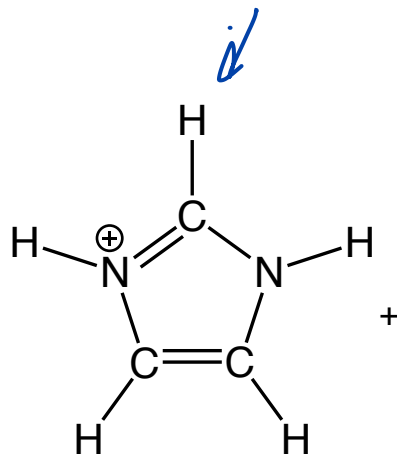
Section 2.12  
Noncovalent Interactions Between Molecules

Chapter 3

**(16) Second Class from Today**

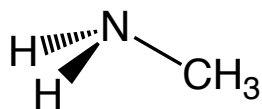
**Third Class from Today (17)**

BL acid proton donor



imidazole

+

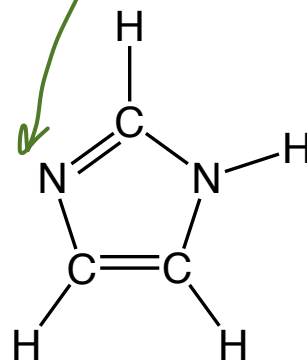


BL base

methylamine

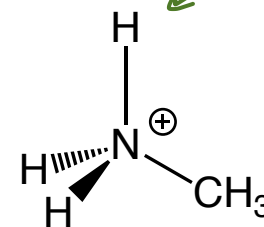


accepts the proton



the H is gone

+



here it is

Will it go?  
yes, the protonated imidazole ring will donate it's proton to the methyl amine

$K_a$  and  $pK_a$

an acid dissociation equilibrium constant



$$pH = 7$$

$$[H^+] = 1 \times 10^{-7} = 0.0000001$$

pH?

$$pH = -\log[H^+]$$

$$K_a = \frac{[products]}{[reactants]}$$

.1 ..... .00000  
.0000000001

$$K_a = \frac{[H^+][A^-]}{[HA]} = \text{small?} = \frac{\text{small}}{\text{big}} \frac{1}{4}$$

a strong acid has a large or small  $K_a$ ? Big

Large  $K_a$  means lots of products  
few reactions  $\frac{10000}{1}$

the larger the  $K_a$  the stronger the acid

$$pK_a = -\log K_a$$

$$pK_a = -\log K_a$$

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

$$-\log K_a = -\log \left( \frac{[H^+][A^-]}{[HA]} \right)$$

rearrange

$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

Why Tables of  $pK_a$  instead of  $K_a$ ? Buffers

minimize changes in pH  
when acids or bases are added  
to the solution

Section 2.8

$pK_a$  of acetic acid is 4.74

$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

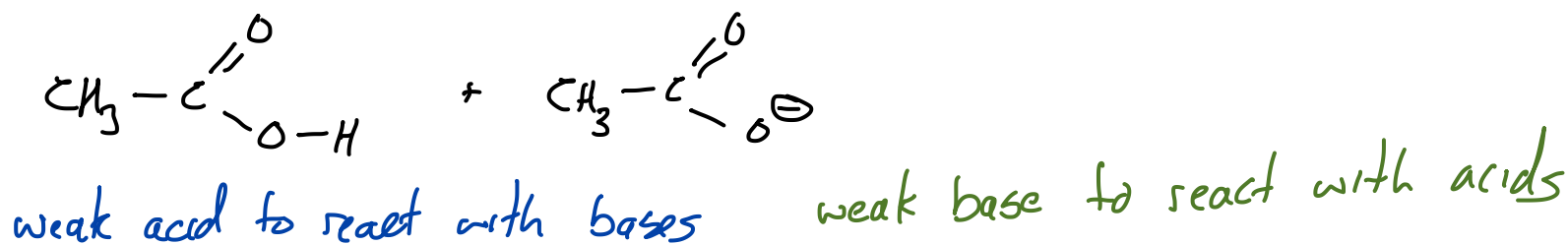
Buffer has to be a base to grab onto  $H^+$ 's  
that are added

Buffer has to be an acid to grab the base that  
was added

Buffer must be a mix of an acid + base



Mixtures of weak acids + weak bases make buffers



Why Tables of  $pK_a$  instead of  $K_a$ ? Buffers

minimize changes in pH  
when acids or bases are added  
to the solution

Section 2.8

$pK_a$  of acetic acid is 4.74

$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

Acetate buffer with a pH of 5?

$$5.00 = 4.74 + \log \frac{[A^-]}{[HA]}$$

$$0.26 = \log \frac{[A^-]}{[HA]}$$

$$10^{0.26} = \frac{[A^-]}{[HA]}$$

so now I know how much  
 $A^-$  I need for  
amount of HA I  
choose



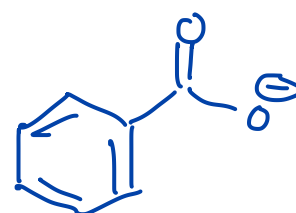
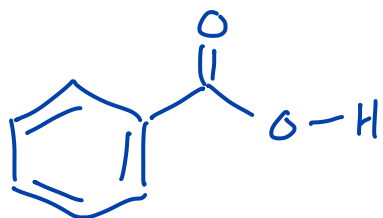
$$K_a = 10^{-4.74}$$

Why Tables of  $pK_a$  instead of  $K_a$ ? Solubility

~~acetic acid~~  
benzoic acid  
 $pK_a$  of ~~acetic acid~~ is 4.20

the HH equation at a given pH tells us which species dominates

$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$



not particularly water soluble

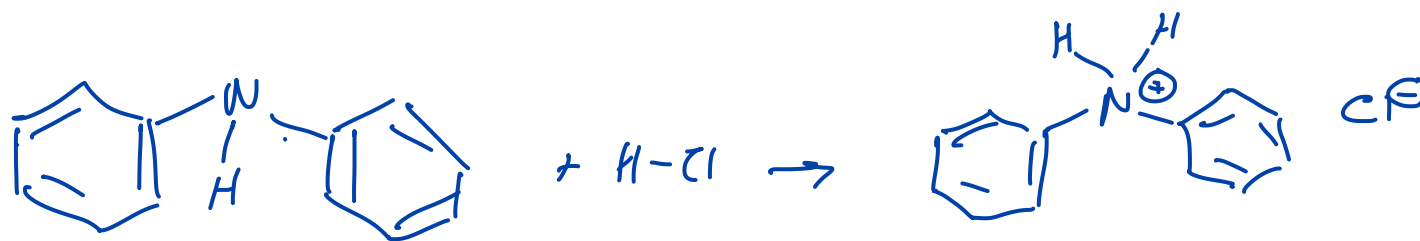
much more soluble in organic solvents

much more water soluble

$$pH = 4.2 + \log \frac{[A^-]}{[HA]} \quad \frac{100}{1}$$

$$pH = 4.2 + 2$$

$$pH = 6.2$$



Less Soluble in Water  
More Soluble in Organic Solvents

Less Soluble in Organic Solvents  
Increased Solubility in Water



pK<sub>a</sub> - Which is the strongest acid?

Section 2.8

acid? no



NH<sub>3</sub>

smaller/more negative pK<sub>a</sub>  
is the stronger acid

H<sub>2</sub>O

$\text{p}K_a = 3.18$

$K_a = 10^{-3.18}$  stronger

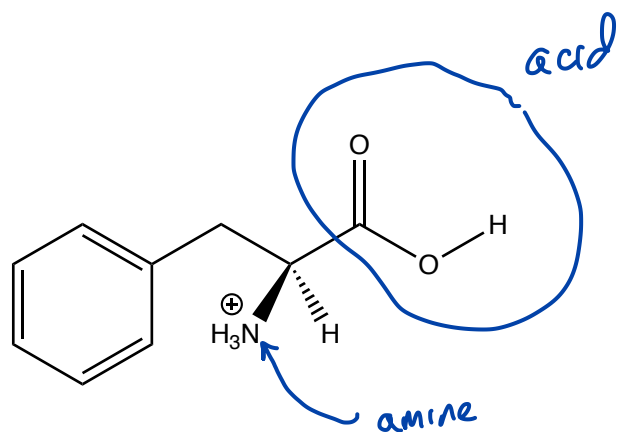


$-\log K_a = \text{p}K_a$

pK<sub>a</sub>'s    CH<sub>4</sub>, ~50    NH<sub>3</sub>, ~36    H<sub>2</sub>O, 15.6    HF, 3.18

pK<sub>a</sub> - Which is the stronger acid?

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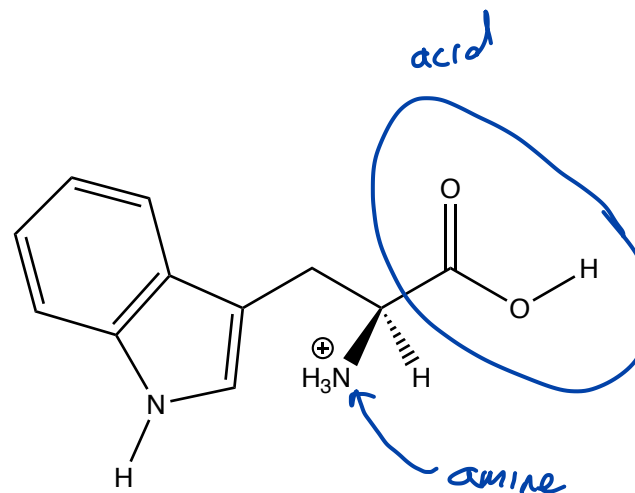


phenylalanine

pK<sub>a</sub> = 1.83

stronger acid

$$K_a 10^{-1.83}$$



tryptophan

pK<sub>a</sub> = 2.83

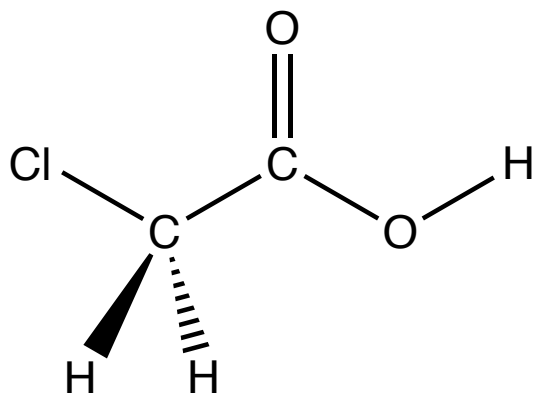
ammonium ion

$$K_a 10^{-2.83}$$



pK<sub>a</sub> - Which is the stronger acid?

Section 2.8



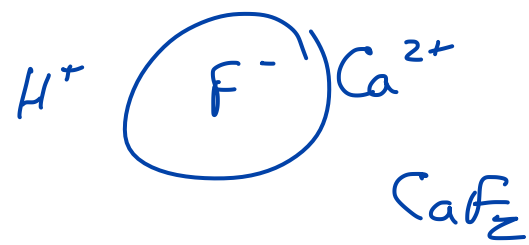
pK<sub>a</sub> = 2.87

stronger



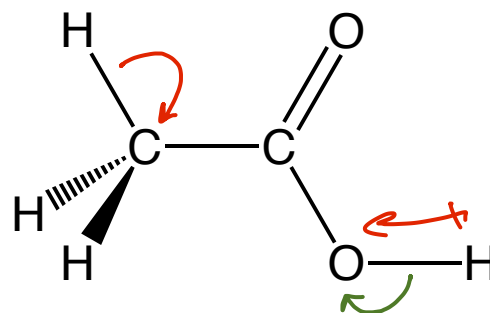
3.18  
pK<sub>a</sub> = ~~2.87~~

weak but very dangerous



Which is the acidic H<sup>+</sup> in acetic acid (HC<sub>2</sub>H<sub>3</sub>O<sub>2</sub>)?

Section 2.6



a.k.a. CH<sub>3</sub>CO<sub>2</sub>H

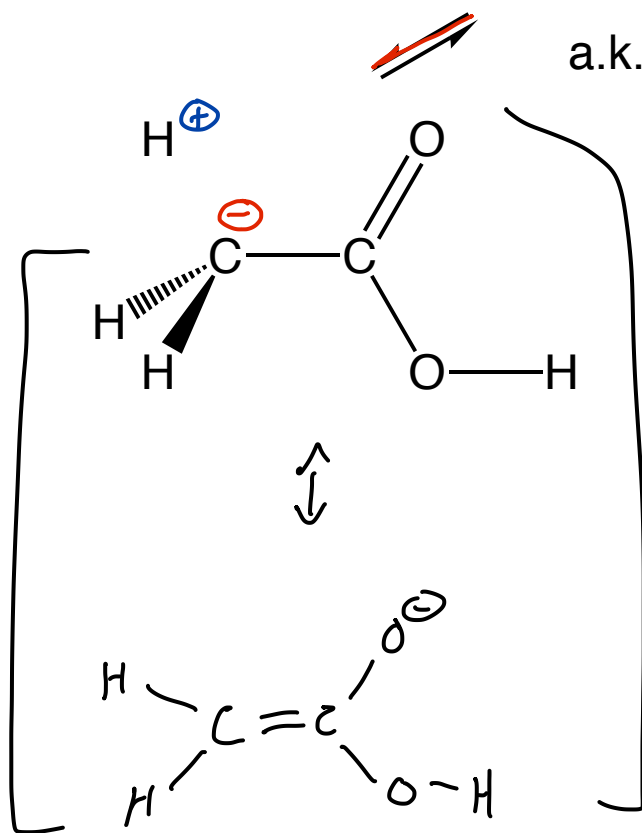
O atom has 8 ip<sup>+</sup> in its nucleus and is betterable to support the ⊖ charge

The atom that supports the ⊖ charge

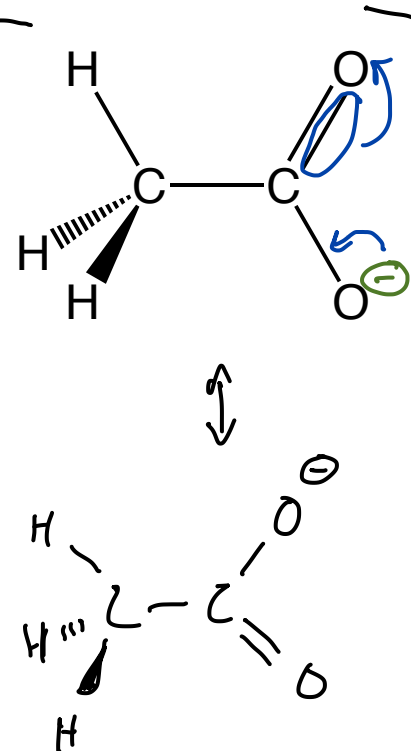
better is the one where

H<sup>+</sup> the bond

to the H<sup>+</sup> will break



?



The one that leaves the more/most stable base behind