

**(11) Today**

Sections 2.7 – 2.11: Acids and Bases

Section 2-12: Non-Covalent Interactions  
Between Molecules

Section 2.7: Problem 2-11

Section 2.8: Problems: 2-12, 2-13

Section 2.9: Problems 2-14 – 2-16

Section 2.10 - 2:11: 2.11 Problems: 2-17,  
2-18, 2-24, 2-25, 2-40 (2-42 is a good  
question but the Lewis acid-base concept is  
not strongly emphasized in our organic class),  
2-43, 2-44, 2-46, 2-47, 2-48, 2-54, 2-55, 2-61,  
2-64

**Next Class (12)**

Section 2-12  
Non-Covalent Interactions Between  
Molecules

2.12 Problems: 2-19, 2-65

**(13) Second Class from Today**

Section 3.1  
Functional Groups

Section 3.2  
Alkanes and Isomers

Section 3.3  
Alkyl Groups

Section 3.4  
Nomenclature

**Third Class from Today (14)**

Section 3.1  
Functional Groups

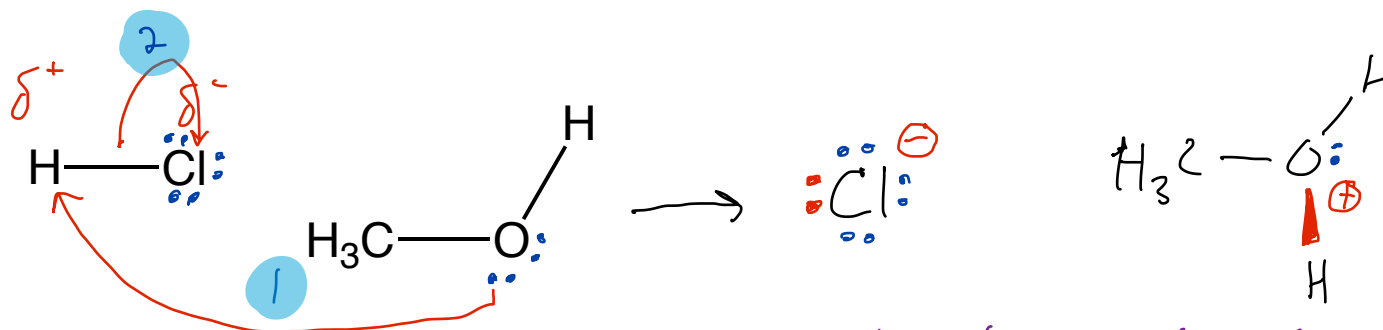
Section 3.2  
Alkanes and Isomers

Section 3.3  
Alkyl Groups

Section 3.4  
Nomenclature

A Brønsted-Lowry acid is a proton,  $H^+$ , DONOR.

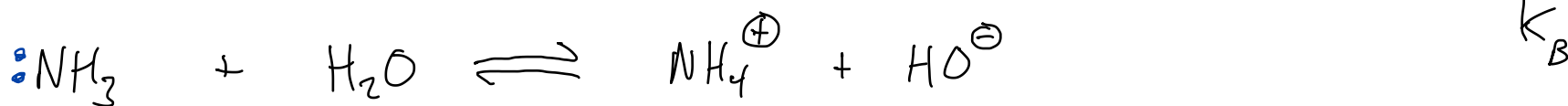
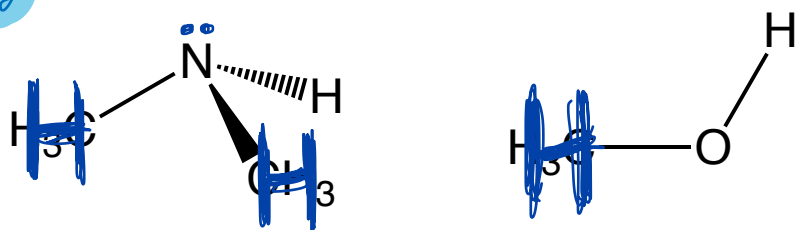
A Brønsted-Lowry base is a proton,  $H^+$ , ACCEPTOR.

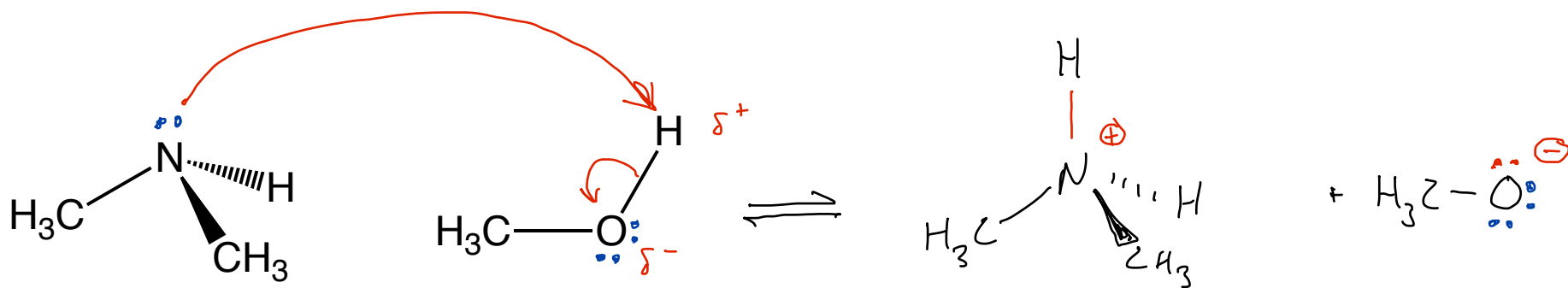
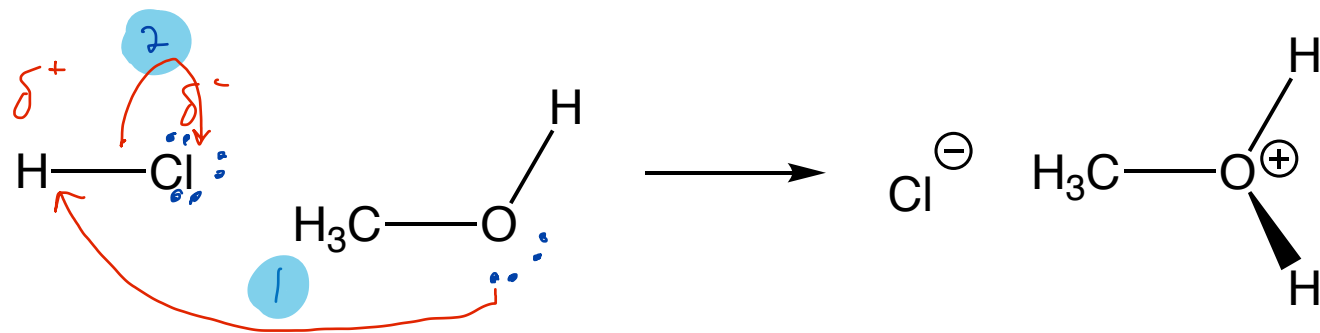


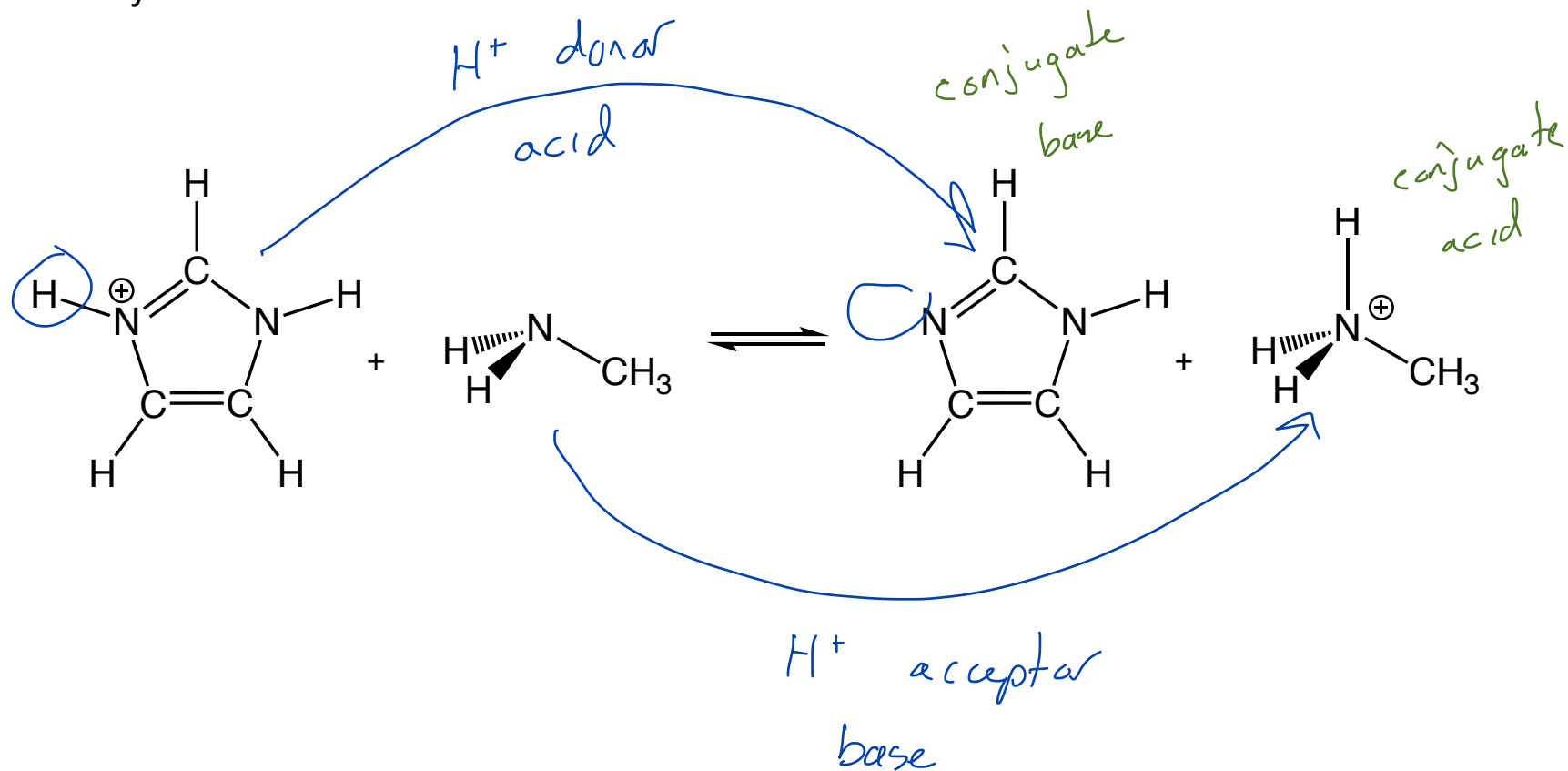
Curved arrow notation: a way to keep track of where we are making + breaking bonds

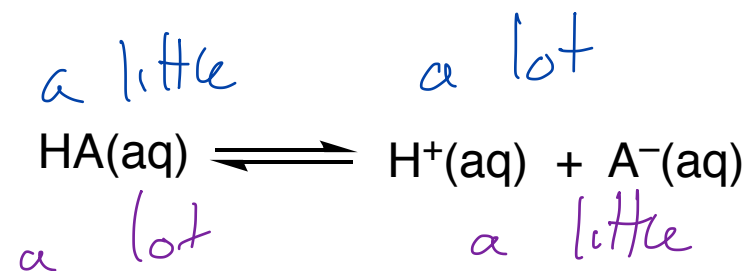
arrows start at  $e^-$ 's and the arrow ends at where the  $e^-$ 's are going

- 1 in this case the arrow means we are making an O to H bond
- 2 in this case the arrow means the  $e^-$ 's in the bond are moved to the Cl









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

a lot
a little

Do strong acids (things that are good at releasing H<sup>+</sup>) have **large** or small K<sub>a</sub>'s?

weak acid  $\left( \begin{array}{l} \text{small} \\ \text{big} \end{array} \frac{\text{a little}}{\text{a lot}} = K_a \right)$  small K<sub>a</sub>

the "p" in pK<sub>a</sub> is the same p as the p in pH

$$pK_a = -\log K_a$$

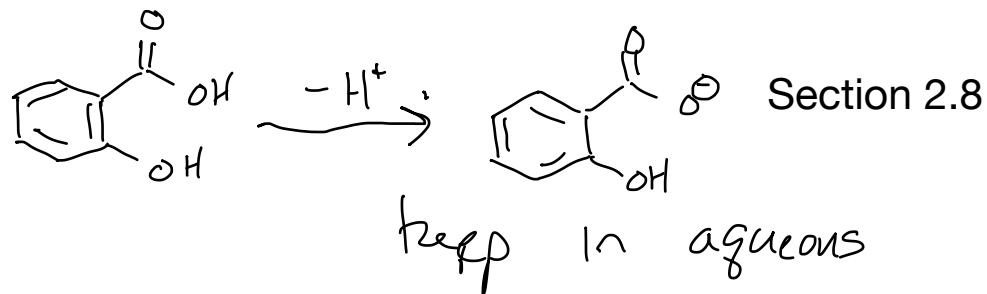
$$K_a = \frac{[H^+][A^-]}{[HA]}$$

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

rearrange

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

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



acetate buffer  
@  $pH = 4.00$

$$pK_a = -\log K_a$$

buffers

sodium acetate

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

acetate / acetic acid

$$4.00 = 4.78 + \log \frac{[\text{acetate}]}{[\text{acetic acid}]}$$

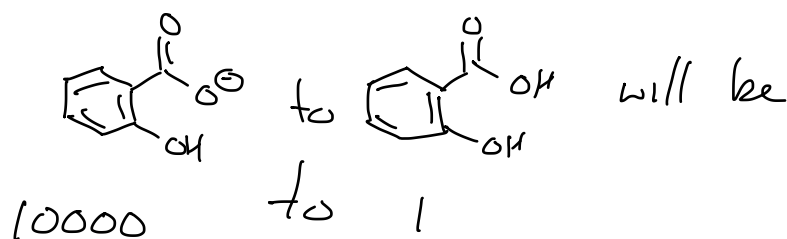
$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

10000 / 1

$$= 2.98 + \log 10000$$

$$pH = 2.98 + 5$$

at a pH of 8 the





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

$$pK_a = -\log K_a$$

Section 2.8

pK<sub>a</sub>

50

CH<sub>4</sub>

$$K_a = 10^{-50}$$

the larger the pK<sub>a</sub>  
the weaker the  
acid

36

NH<sub>3</sub>

15.6

H<sub>2</sub>O

3.18

HF

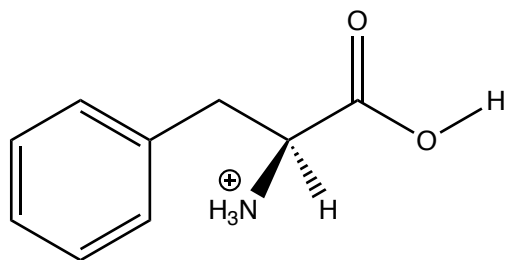
strongest acid

the smaller the pK<sub>a</sub>  
the stronger the  
acid

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

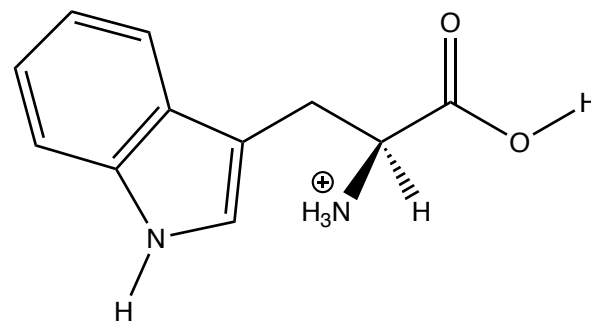
pK<sub>a</sub> - Which is the stronger acid?

Section 2.8



phenylalanine

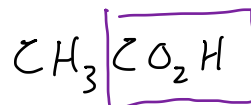
pK<sub>a</sub> = 1.83



tryptophan

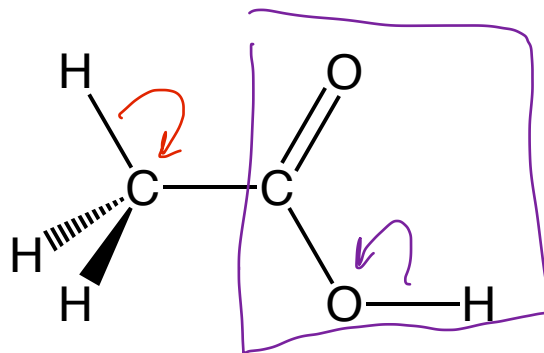
pK<sub>a</sub> = 2.83

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



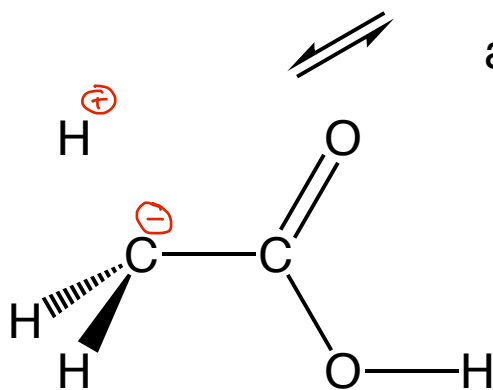
carboxylic acid

Section 2.6

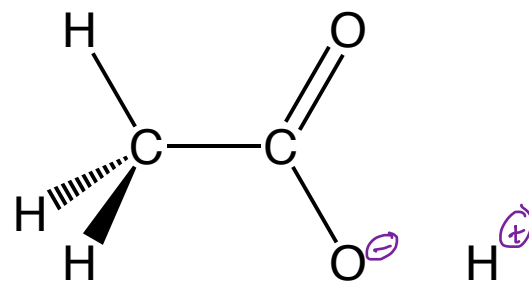


a.k.a. CH<sub>3</sub>CO<sub>2</sub>H

this side



?



C atoms have a 6<sup>+</sup> nucleus  
 these e<sup>-</sup>'s experience less attraction  
 these e<sup>-</sup>'s would be less stable  
 these e<sup>-</sup>'s would be higher in E

O atoms are better at holding onto e<sup>-</sup>'s...  
 O atoms have 8<sup>+</sup> nucleus  
 e<sup>-</sup> experience a greater ⊕ charge... more attraction...  
 lower in E

The one that leaves the more/most stable base behind

Five ways to stabilize the electrons on the conjugate base

Section 2.8, 2.10

Same Period More Positive Nucleus

CH<sub>4</sub>

NH<sub>3</sub>

H<sub>2</sub>O

HF

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

Five ways to stabilize the electrons on the conjugate base

Section 2.6 – 2.9

Same Column Larger Valence Shell

HI

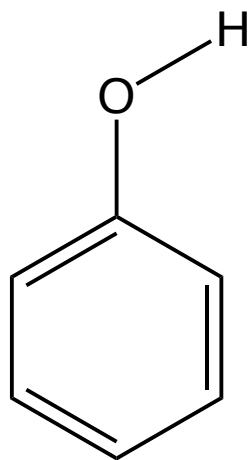
HBr

HCl

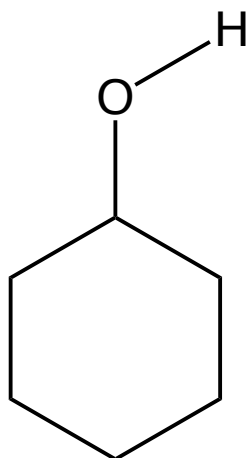
HF

pK<sub>a</sub>'s HF, 3.18 HCl, -7 HBr, -9 HI, -10

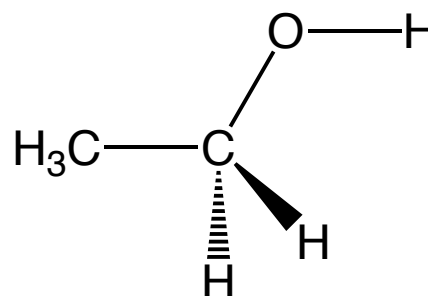
Resonance



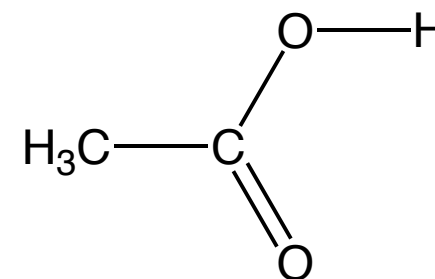
phenol



cyclohexanol



ethanol



acetic acid

pK<sub>a</sub>'s    cyclohexanol, 16.0    phenol, 10.0    ethanol 16.0    acetic acid 4.74

Five ways to stabilize the electrons on the conjugate base

Section 2.6 – 2.9

## Inductive Effect

acetic, 4.76; formic, 3.75; chloroacetic, 2.87; dichloroacetic, 1.25

Five ways to stabilize the electrons on the conjugate base

Section 2.8, 2.10

Greater s character

$pK_a$ 's ethane 50, ethene 44, ethyne 25



Practice: For each molecule, which proton is the most likely to be lost and for each pair, which is the stronger acid

