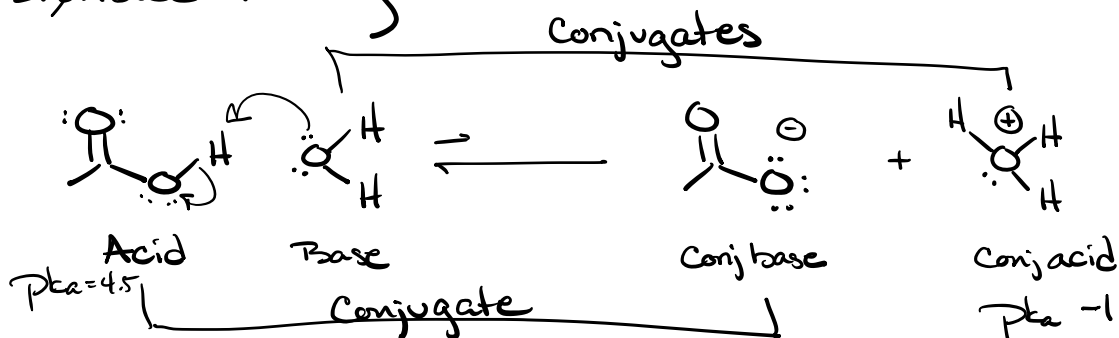


# Acid - Base

Brønsted-Lowry



Acid = proton donor

Base = proton acceptor

At equilibrium the weaker acid & weaker base are the major species in solution.

Strong Acid  $\rightarrow$  Conj. Base of Strong Acid  
= Weak Base

Increasing strength of Acid  $\uparrow$

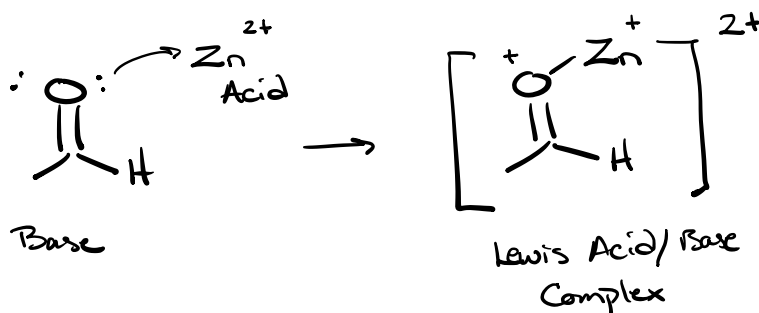
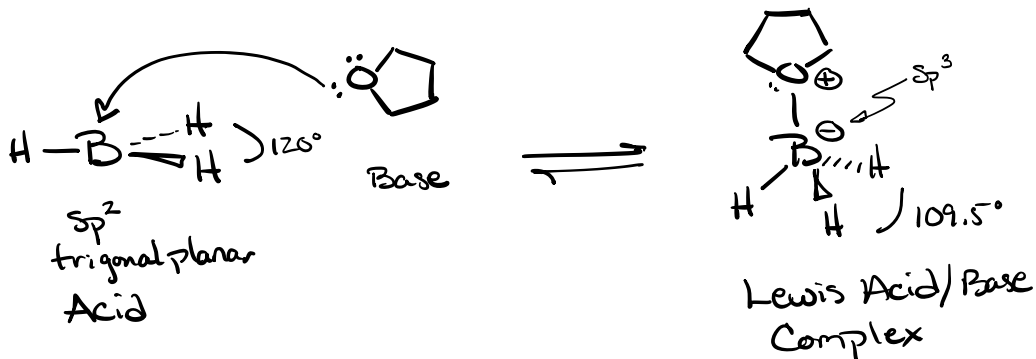
Increasing strength of base  $\downarrow$

Weak Acid  $\rightarrow$  Conj Base of weak acid  
= Strong Base

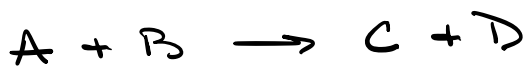
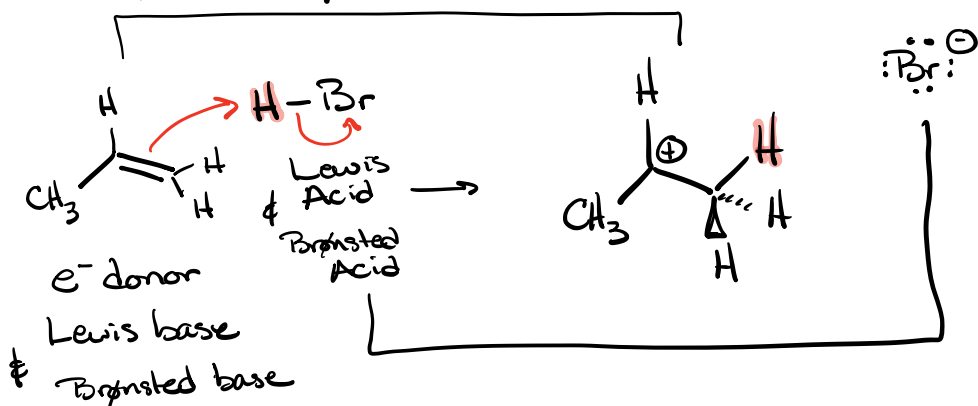
# Lewis Acid-Base (focus on e<sup>-</sup>)

Lewis Acid - e<sup>-</sup> acceptor

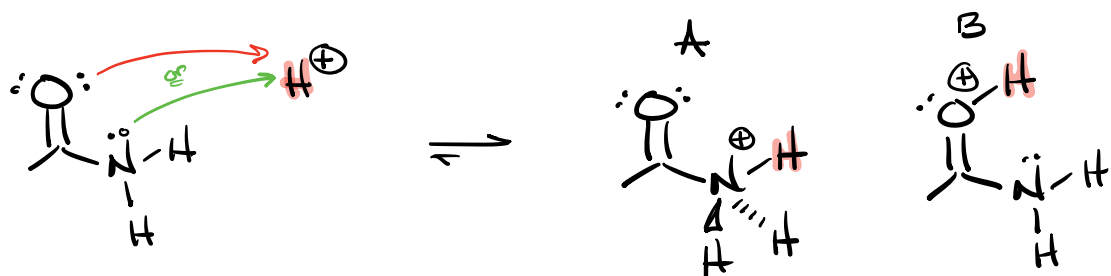
Lewis Base - e<sup>-</sup> donor



Compounds w/ π-bonds can also be Lewis base



+ = "And"



which lone pair is more basic?

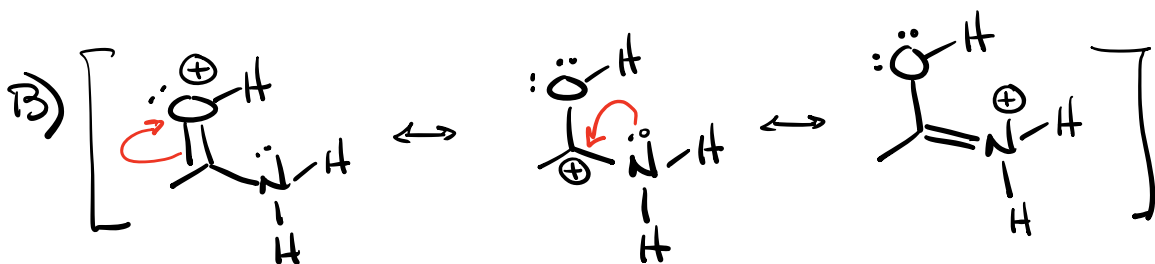
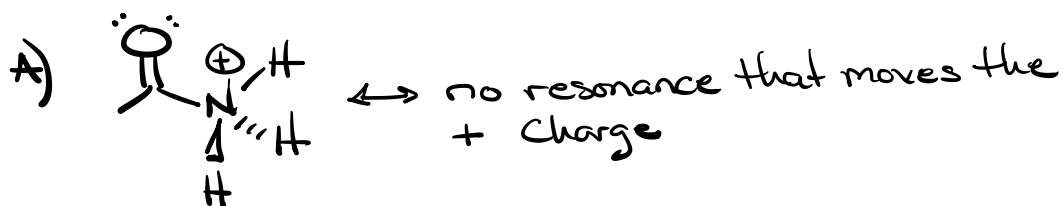
which location is more likely to be protonated?

we can ask which product is more stable?  
(which is lower in energy?)

more stable = less reactive = weaker Acid  
or  
weaker Base } As appropriate

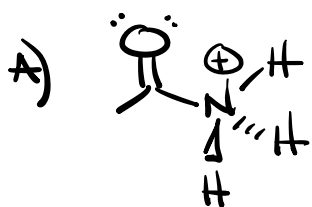
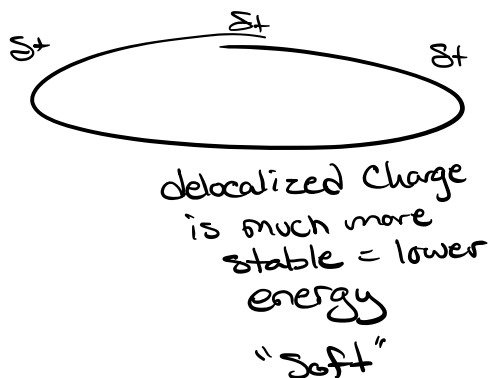
How do we assess relative stability?

⇒ Resonance for one, there are other factors that we will see later.

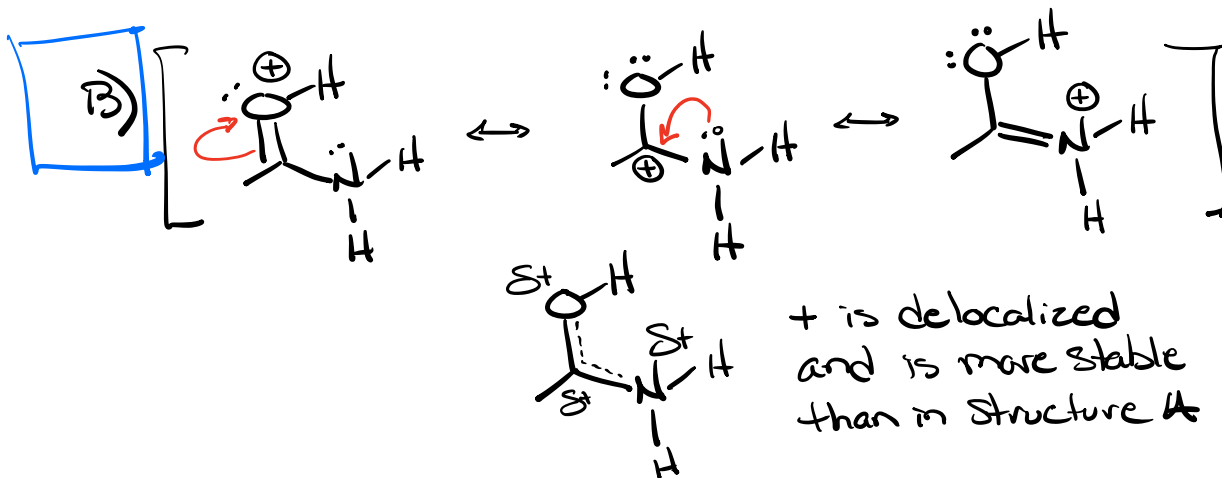


A structure with more resonance contributors, \*that are good resonance contributors, is more stable than one with less. This is due to the delocalization of charge.

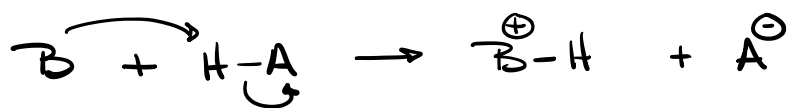
$\text{O}^{\oplus}$   
 Small "Hard"  
 Ion  
 very High in  
 energy



+ charge is localized  
 Small & Hard = High energy  
**Strong Conj. Acid**




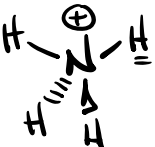
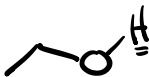
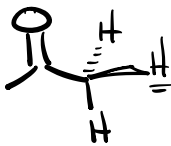
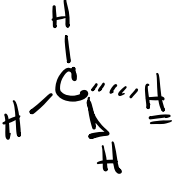
weaker Conj Acid  $\Rightarrow$  more likely to be protonated

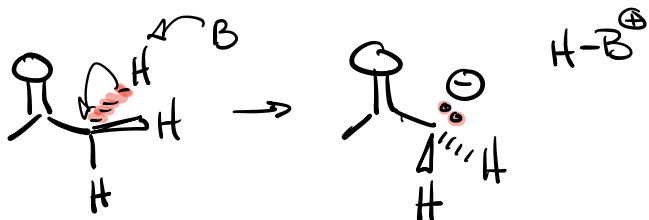
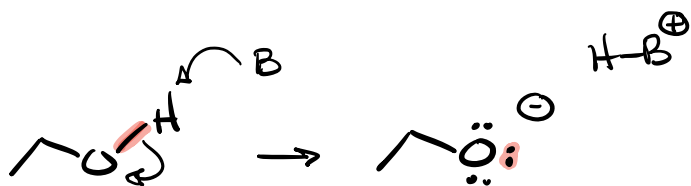


$$K_a = \frac{[B-H^+][A^-]}{[B][H-A]}$$

$$pK_a = -\log K_a$$



|      | <u>pKa</u>               |  | <u>pKa</u> |
|------|--------------------------|--|------------|
| HF   | 3.1                      | H <sub>3</sub> O <sup>+</sup>  | -1.7       |
| HCl  | -3.9                     |  | 4.7        |
| HBr  | -5.8                     |  | 9.3        |
| ⇒ HI | -10.4 <i>most acidic</i> |  | 15.2       |
|      |                          |  | 19         |
|      |                          |  | 60         |



B = Base

# How structure affects Acidity & Basicity

- ① strength of the bond w/ proton
- ② Electronegativity of atoms involved
- ③ Resulting delocalization that has many factors

## Contributing Factors

why is HI most acidic?

|            | <u>pKa</u> |                   |
|------------|------------|-------------------|
| $\sim 6$ ↘ | HF 3.1     | F <sup>-</sup> ○  |
| $\sim 2$ ↘ | HCl -3.9   | Cl <sup>-</sup> ○ |
| $\sim 4$ ↘ | HBr -5.8   | Br <sup>-</sup> ○ |
|            | HI -10.4   | I <sup>-</sup> ○  |

2~6 pKa unit difference

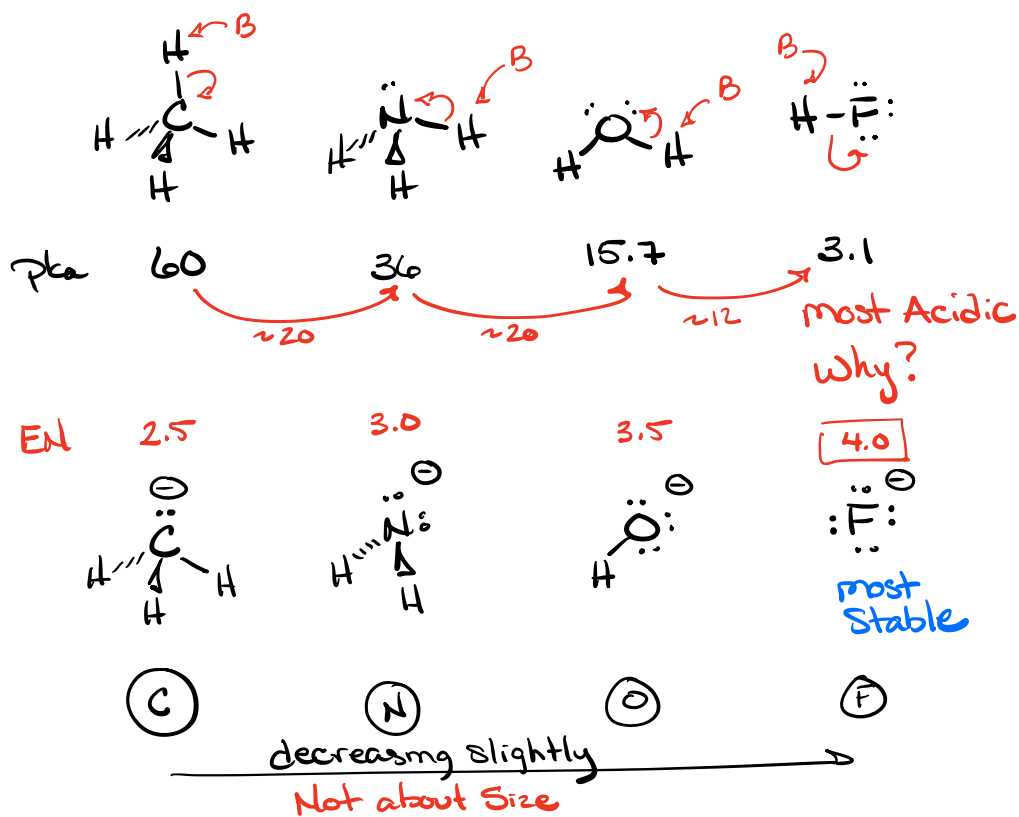
Size!

F<sup>⊖</sup> is most EN  
⇒ should be most stable with negative charge, but it's not!

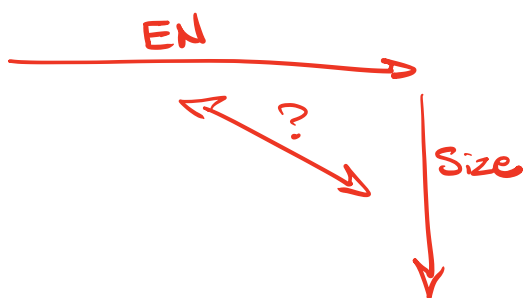
I<sup>⊖</sup> is most stable!!

The more delocalized a charge, the more stable a charge becomes.

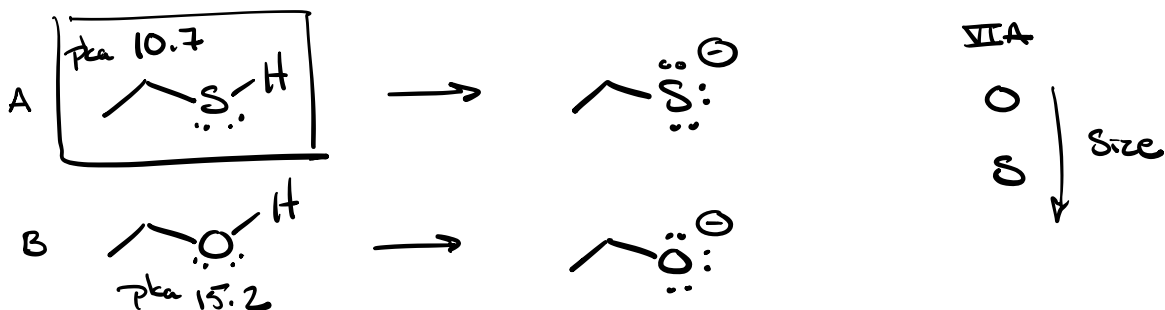
Size > EN ! when going down a column in periodic table



EN is a more important factor when going  
 across the periodic table



which of these is more acidic?

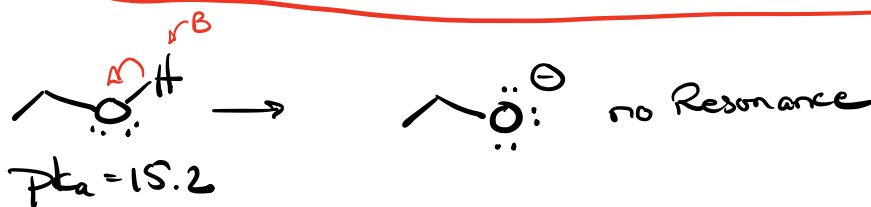
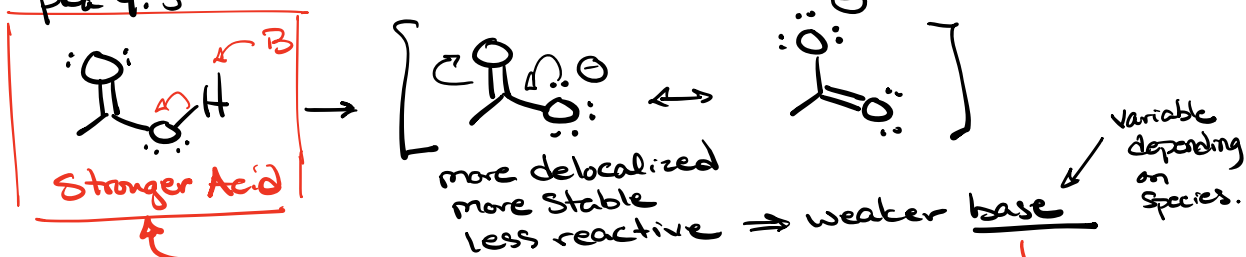


Because Sulfur is larger than oxygen, the Sulfide ion is larger and more delocalized. The sulfide is thus more stable = less reactive = stronger Conj acid!

### Factors

EN 12-20 pKa units  
Size 2-6 pKa units  
Resonance ?  $\sim 10$  pKa units  
Ranking Factors?

which is more acidic?  $\Rightarrow$  look at charged species





One more factor to look at

Induction - the pulling (or delocalization)  
of  $e^-$  through EN differences