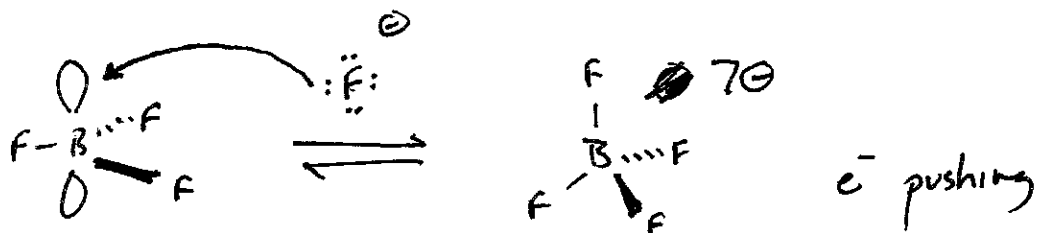


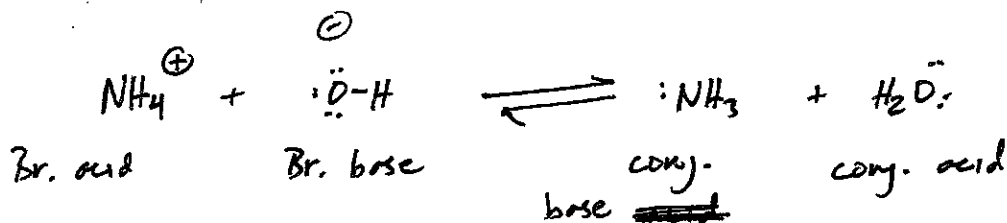
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Last day: - Functional groups + nomenclature
 - introduction into acid/base chemistry (Arrhenius, Bronsted, Lewis)

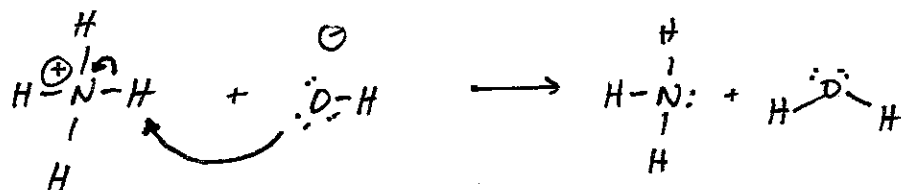
Recall:



Bronsted-Lowry - acid/base chemistry



e⁻ pushing diagram:



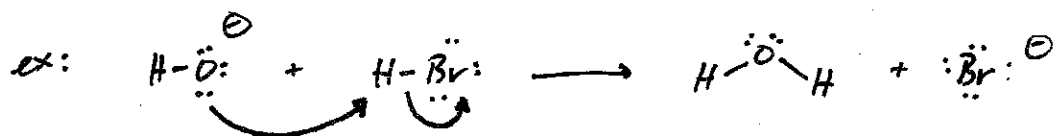
General features:

- 1) Convert acid/base pair to a different acid/base pair (i.e. conj. acid/base)
- 2) A Bronsted base is always a Lewis base
 A Bronsted acid is always a Lewis acid, but not all Lewis acids are B/L acids

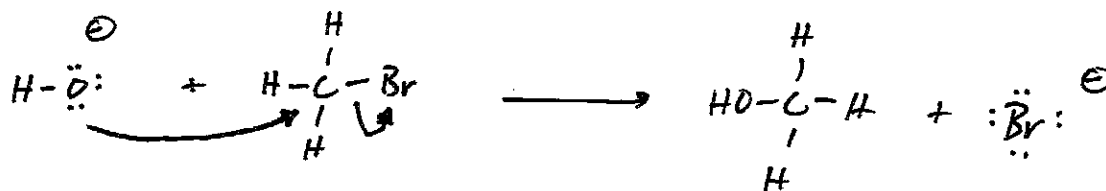
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Relationship between different mechanistic themes

Proton transfer vs. nucleophilic displacement (p 377)



compare to:



bromomethane

- OH^{\ominus} attacks C instead of H
- C gives up one bond ~~(C-Br)~~ (C-Br) in favor of new bond (C-O)

Terminology

Nucleophilic displacement
 $\text{H}-\ddot{\text{O}}:^{\ominus} \rightarrow$ nucleophile
 (nucleus-loving)
 e^- rich
 $\text{H}_3\text{C}-\text{Br} \rightarrow$ electrophile
 (electron-loving)

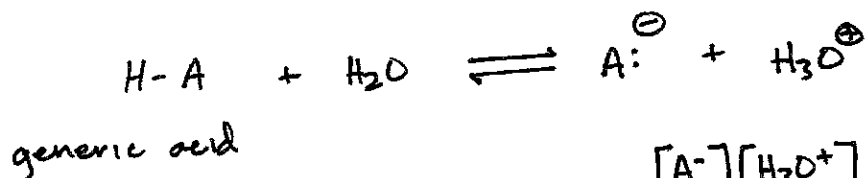
Proton Transfer
 Brønsted base $\rightarrow \text{:}\ddot{\text{O}}-\text{H}^{\ominus}$
 Brønsted acid $\rightarrow \text{H}-\text{Br}$

Br^{\ominus} e^- poor
 leaving group

Conjugate base $\rightarrow \text{Br}^{\ominus}$

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Equilibria and pKa values: Strength of Brønsted acids



$$K_{eq} = \frac{[\text{A}^-][\text{H}_3\text{O}^+]}{[\text{HA}][\text{H}_2\text{O}]}$$

acidity constant (K_a) $K_a = K_{eq}[\text{H}_2\text{O}] = \frac{[\text{A}^-][\text{H}_3\text{O}^+]}{[\text{HA}]}$

for convenience, use log values: $\text{p}K_a = -\log[K_a]$

Table 3A: database of important acids

The stronger the acid,
the lower the $\text{p}K_a$, and
the higher the K_a

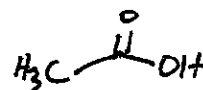
H_2O : $\text{p}K_a$ 16

Alcohol: $\text{p}K_a$ ~16

$\text{H}_3\text{C}-\overset{\text{O}}{\parallel}-\text{OH}$: $\text{p}K_a$ 5

Mineral acids: HCl , $\text{p}K_a$ -6

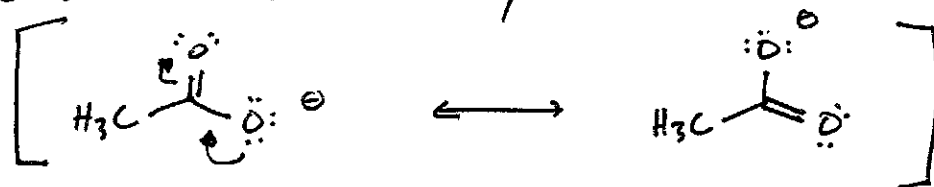
HCl is ~10¹¹ times more acidic than



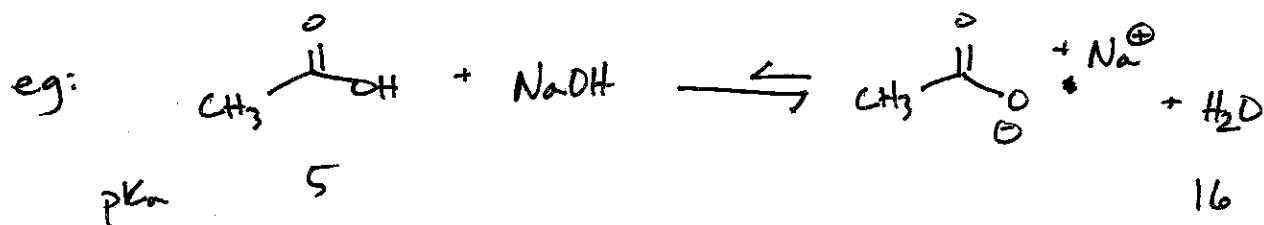
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· Why is ^a ~~a~~ carboxylic acid more acidic than H₂O/alcohol?

- conjugate base is stabilized by resonance



· Knowing characteristic pK_a values allows you to predict positions of acid/base equilibria



equilibria favors weak acid/weak base pairs