

(15) Today

Sections 2.7 – 2.11
Acids and Bases

Lewis Acids

Section 2.12
Noncovalent Interactions Between Molecules

Next Class (16)

Section 3.1
Functional Groups

Section 3.2
Alkanes and Isomers

Section 3.3
Alkyl Groups

Section 3.4
Nomenclature

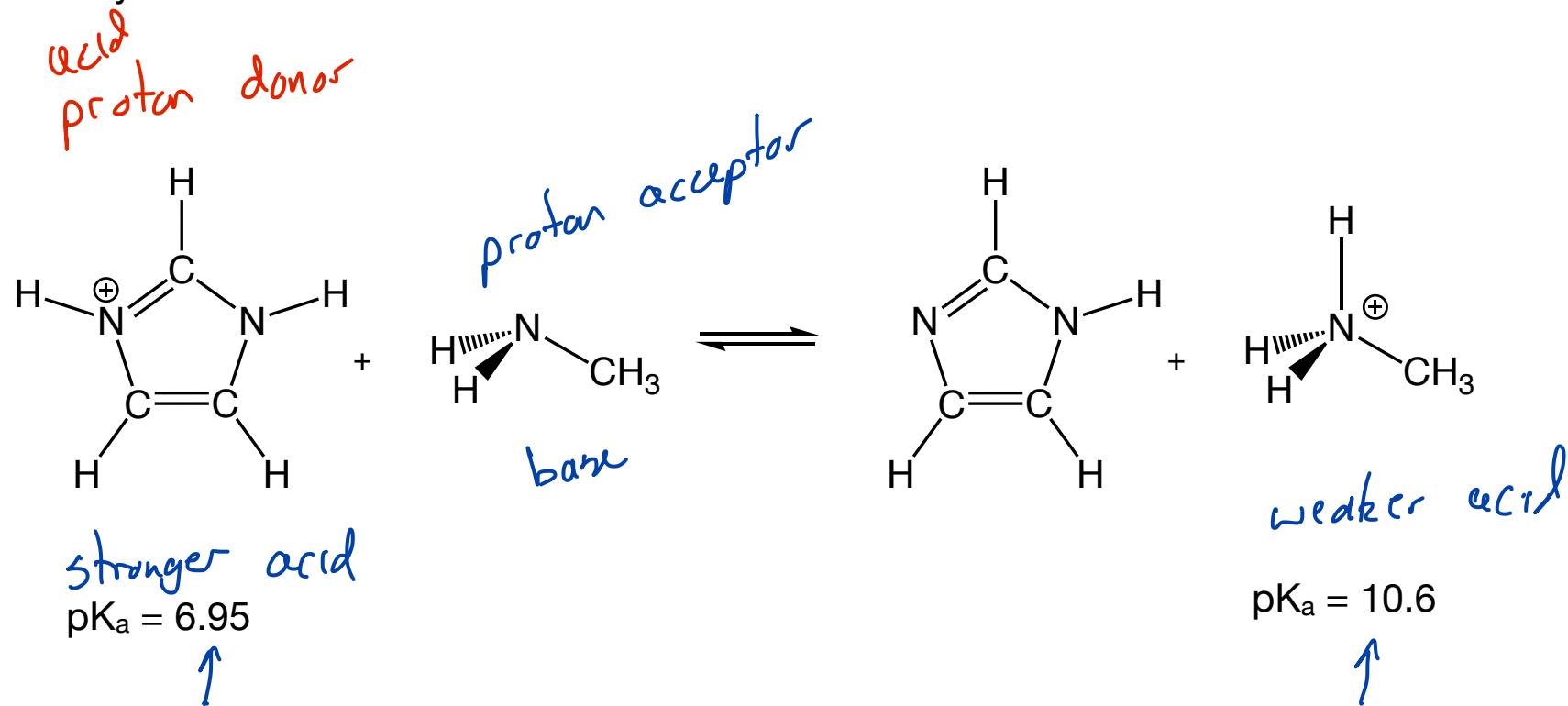
(17) Second Class from Today

Section 3.3 Alkyl Groups
Section 3.4 Nomenclature
Section 3.5 - 3.7 Properties and
Conformations of Alkanes

Third Class from Today (18)

Section 3.5 - 3.7 Properties and
Conformations of Alkanes
Chap 4 Cycloalkanes

nope ... Monday ... for sure



which is the stronger acid...
 the one with a pK_a of 6.95

$$pK_a = -\log K_a$$

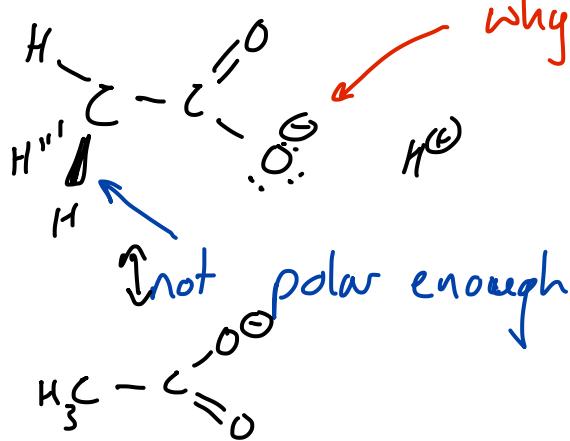
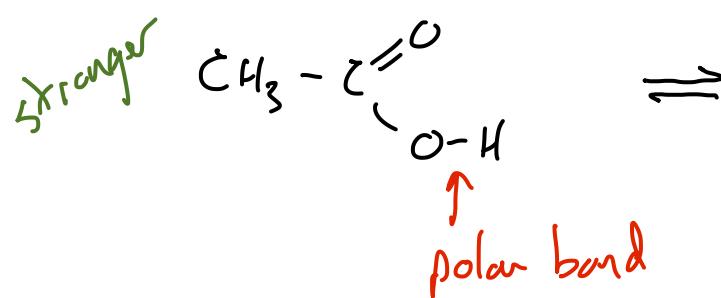
$$K_a \approx 10^{-7}$$

Stronger acid will react
 to give up H^+ and
 make a weaker acid

$$K_a \approx 10^{-11}$$

Acids are H^+ donors the lower the pK_a the stronger the acid.

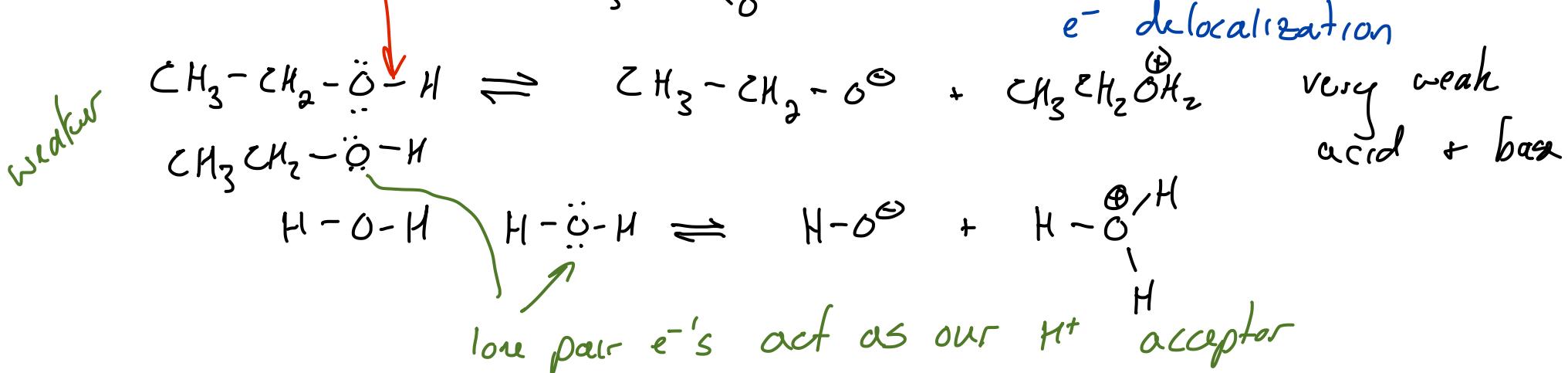
Polar bonds are the bonds most likely to release H^+

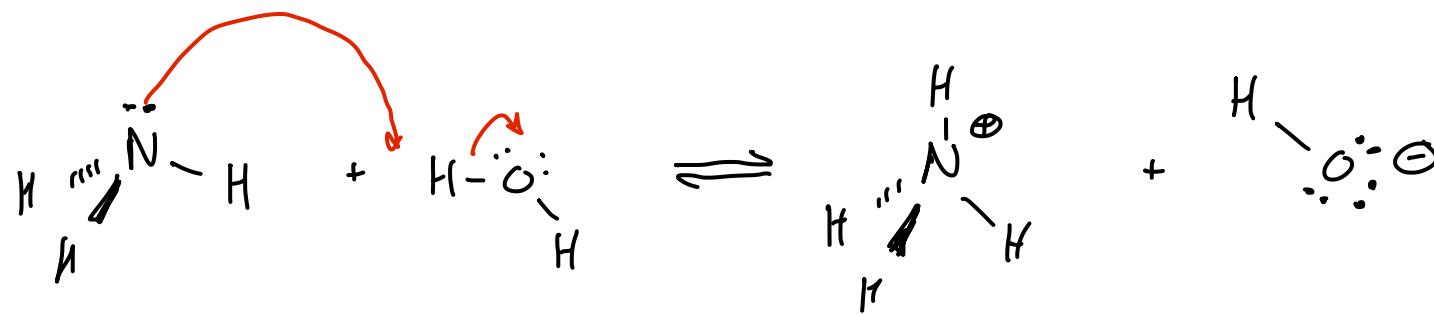


Why are these e⁻'s less reactive

why are these e⁻'s more stable

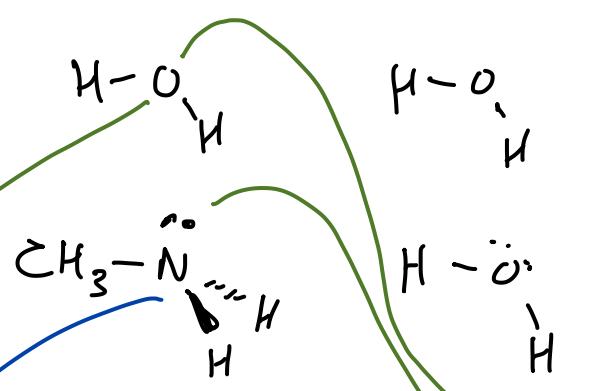
the δ charge is spread out
of 2 O atoms because of
 e^- delocalization



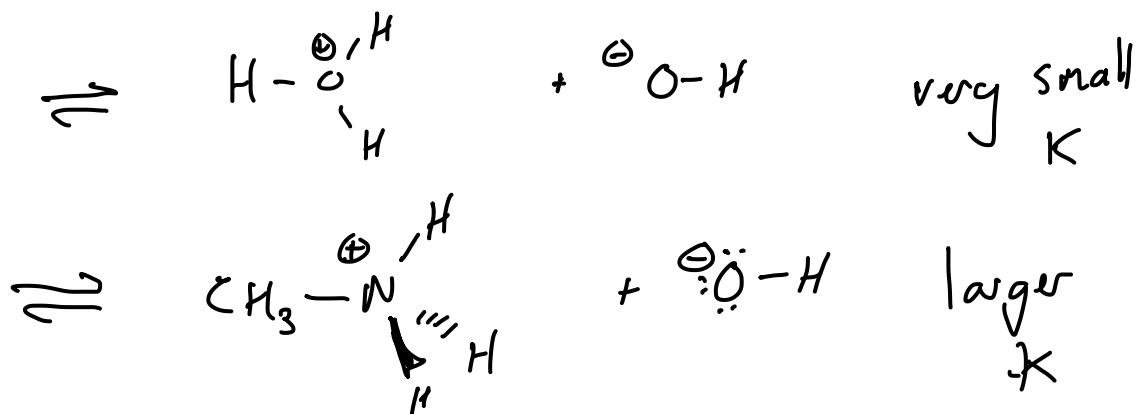


base

acid



Organic amines
are bases



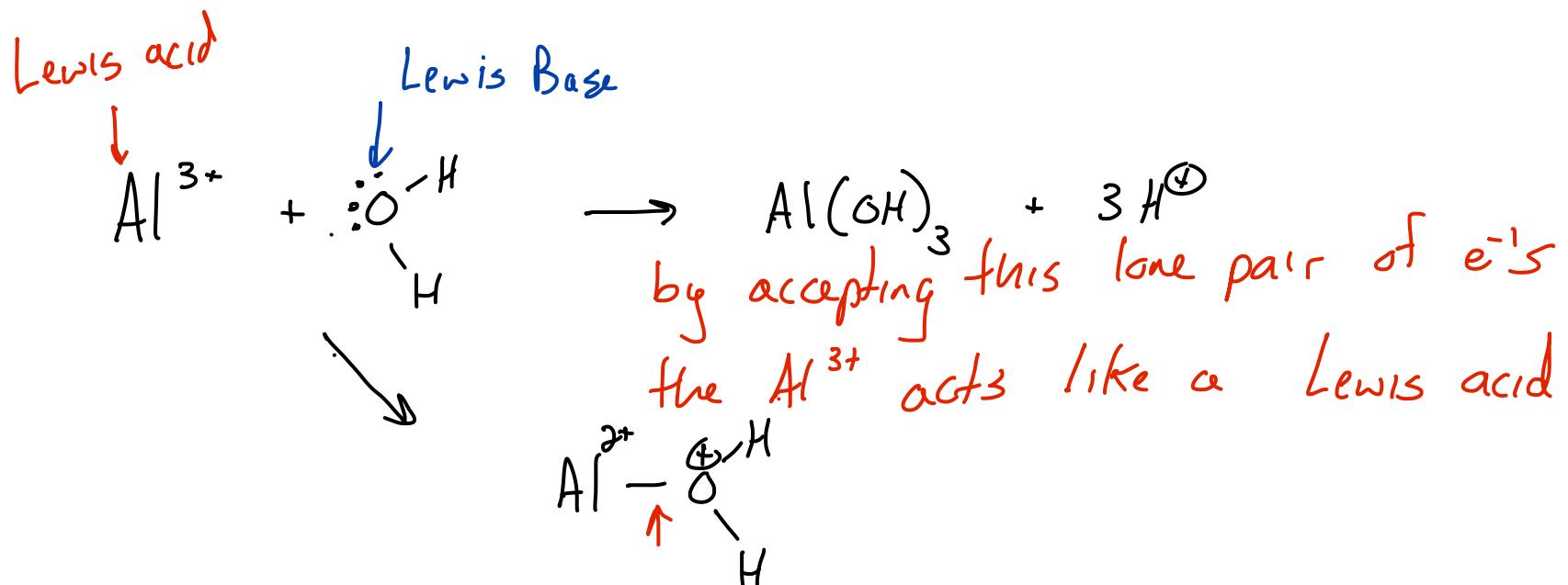
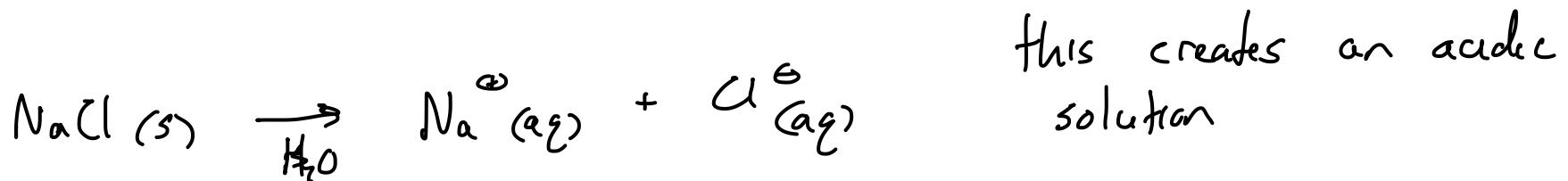
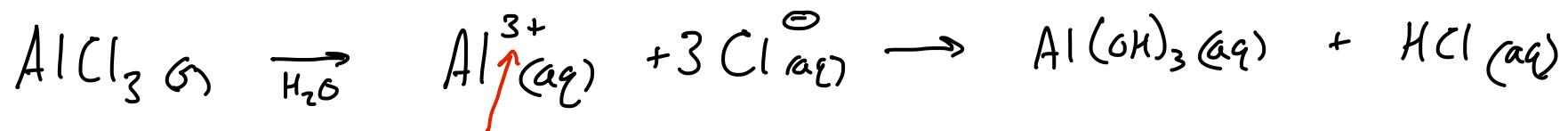
how attractive are these e^- 's

7 ip^+ in N's nucleus.
 e^- 's less stable

N's electrons are less attracted to N's nucleus. Available to react with H^+

8 ip^+ in O's nucleus
 e^- 's are more stable

O's e^- 's are more strongly attracted to O's nucleus. less available to react with H^+



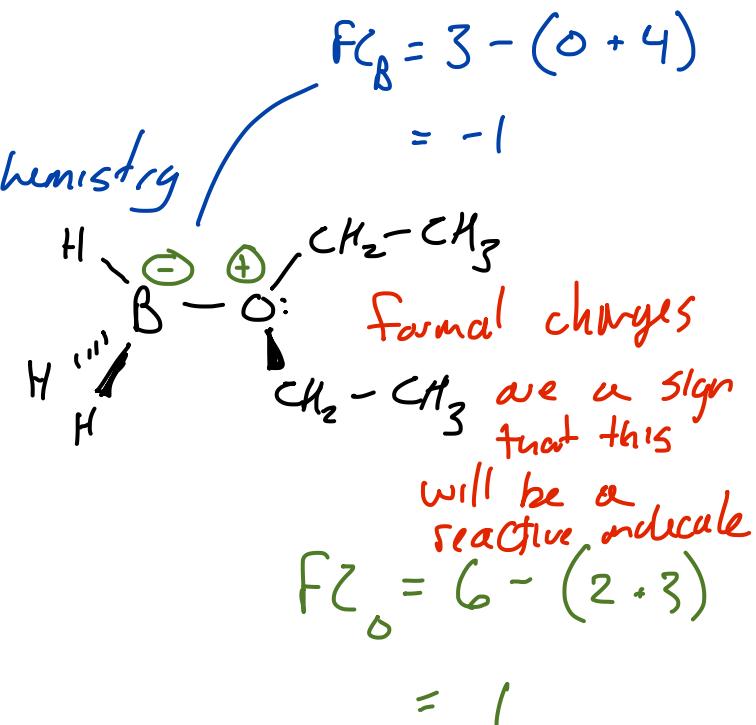
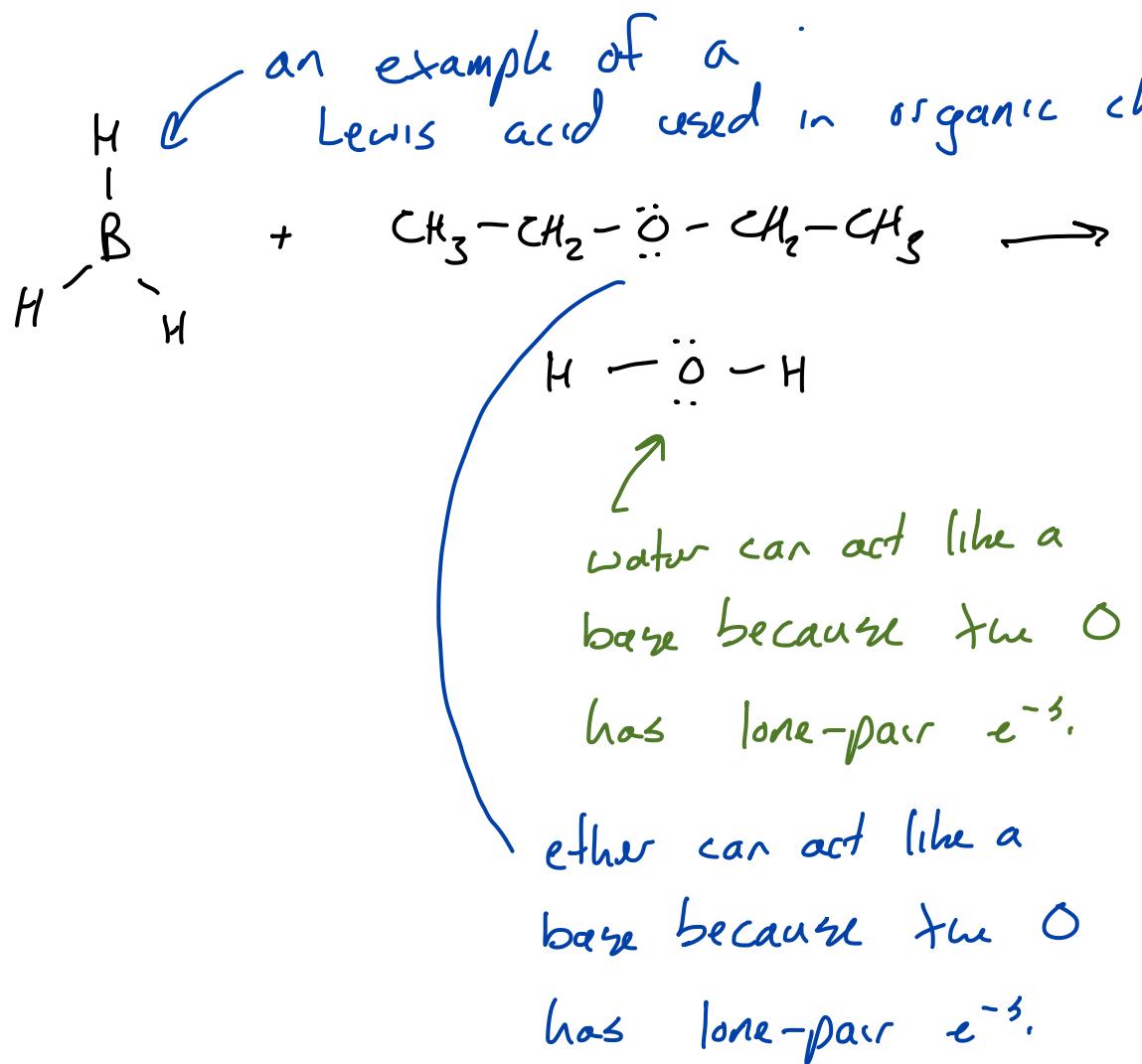
Lewis bases are lone-pair e⁻ donors

metals that are +2 and higher act as Lewis acid





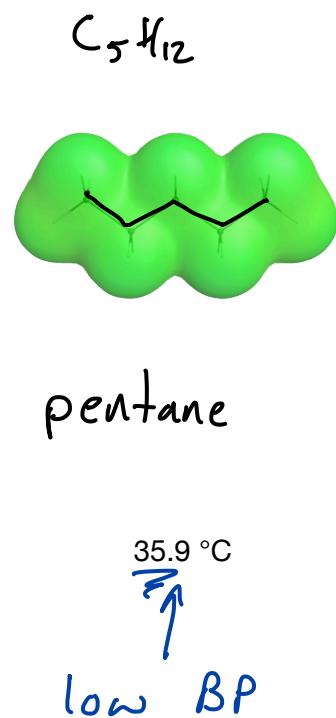
$\text{H} \ddot{\text{B}} \text{H}$ incomplete octet... room for
2 more e^- 's



Oxygen is sharing more e^- 's than we would normally expect

Collectively referred to as...

intermolecular forces, van der Waals forces, or noncovalent interactions



weaker
intermolecular
forces

