

Acid chlorides (acyl chlorides)

R^OCl

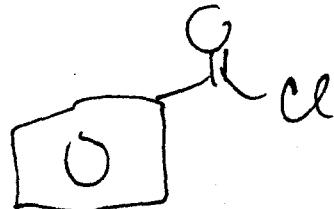
name parent acid

replace "ic acid" with "yl chloride"

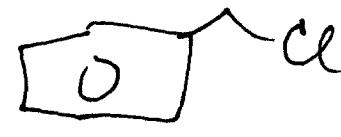
H^OCl acetyl chloride

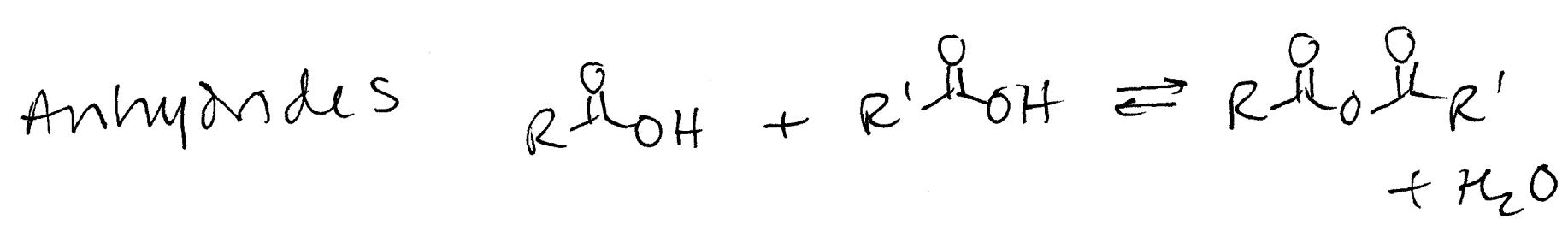


5-methyl-4-hexenoyl chloride



benzoyl chloride

* NOT benzyl chloride → 

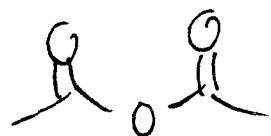


name both parent acids

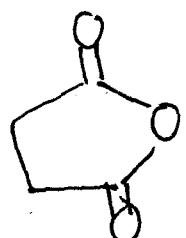
replace "acid" with "anhydride"



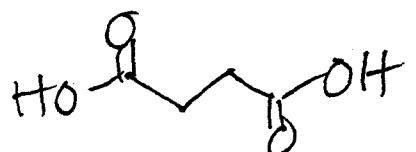
butanoic propanoic anhydride



acetic anhydride

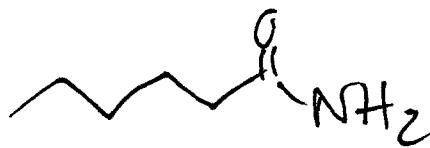


succinic anhydride - from



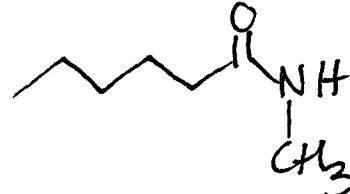
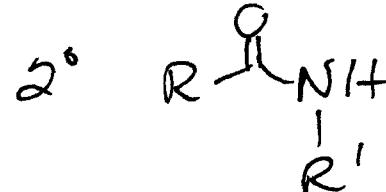
succinic acid

Amides



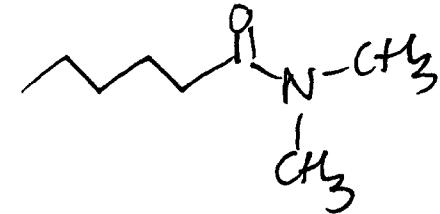
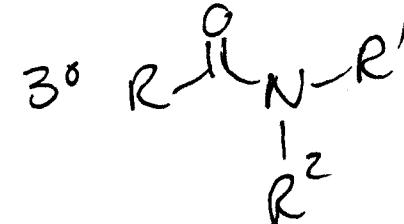
hexanamide

"oic acid"
⇒ "amide"



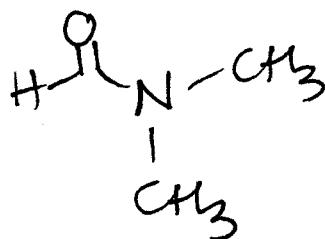
hexanamide

BUT -
what about
the CH_3 ?



DMF

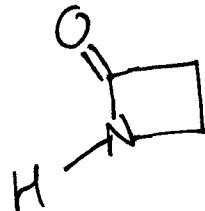
N,N -dimethylformamide



N-methyl hexanamide

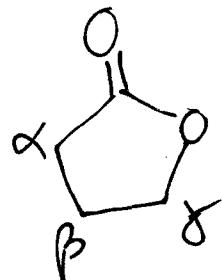
N,N -dimethylhexanamide

cyclic amides are called lactams



a β -lactam

cyclic esters are called lactones



a γ lactone

Nitriles - are considered derivatives of carboxylic acids

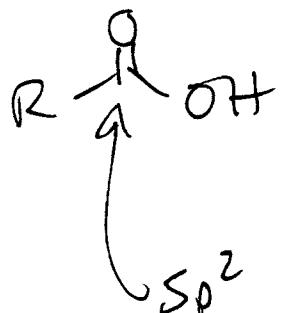


pentano nitrile

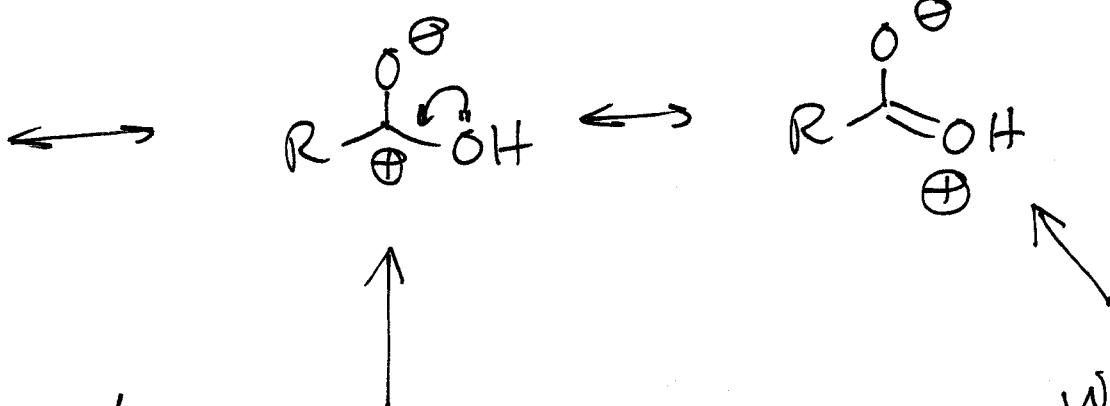


benzonitrile

structural features



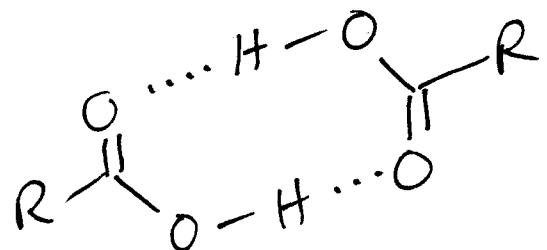
sp^2
120° bond angles



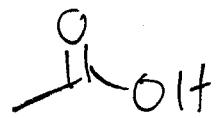
explains
why carbonyls
are susceptible
to nucleophilic attack

explains
why acids
+ derivs
are less
reactive
than ald.
or ketones

carboxylic acids can form H-bond pairs



increases both
mp + bp

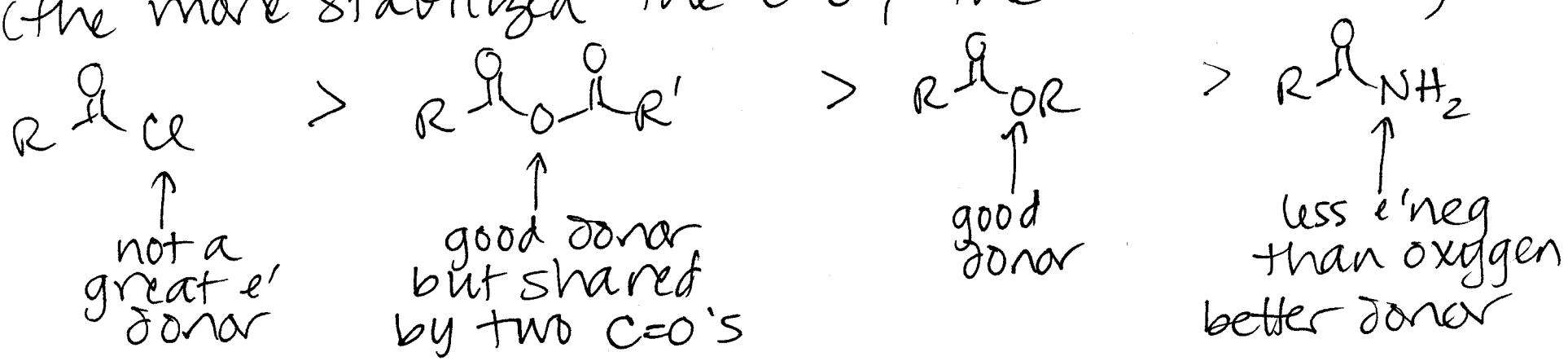


$$\begin{array}{r} 24+4+32 \\ \hline 60 \end{array}$$

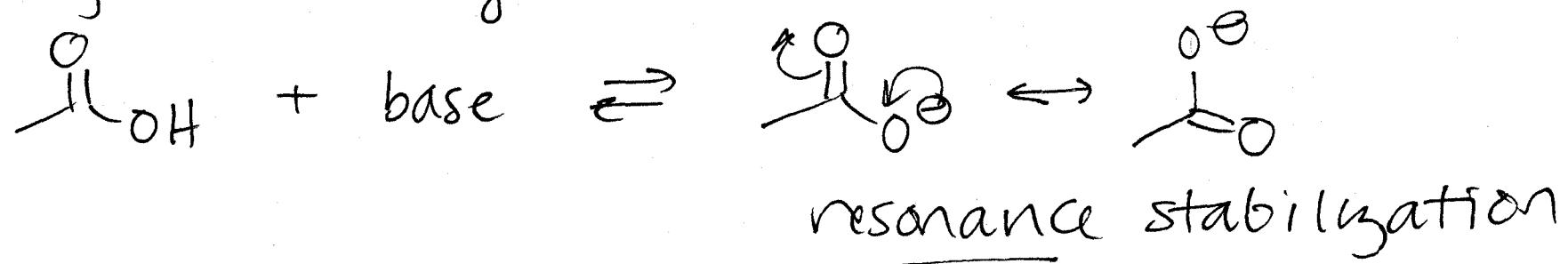
= 60 g/mol

look at derivatives - e' donation from the substituent stabilizes the C=O, making it less susceptible to nucleophilic attack.

(the more stabilized the C=O, the less reactive)



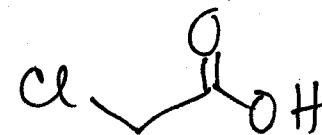
Acidity of carboxylic acids



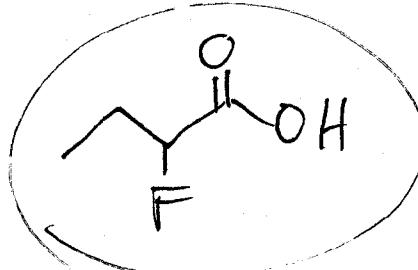
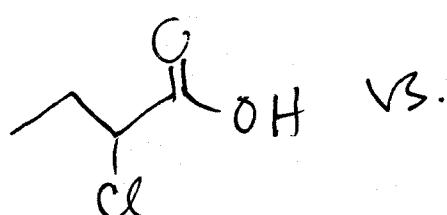
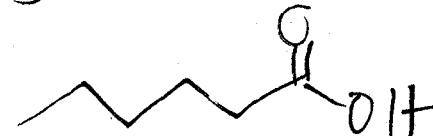
e'neg substituents (e' withdrawers) close to OH will stabilize the corresponding anion by induction. \Rightarrow increased acidity.



pKa 4.7



2.9



vs.

