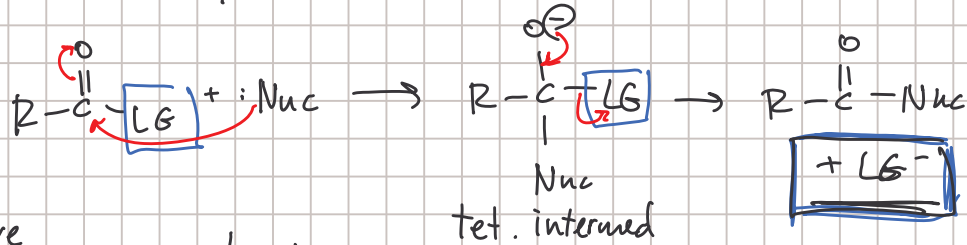


Interconversion of carbox. acid derivs.

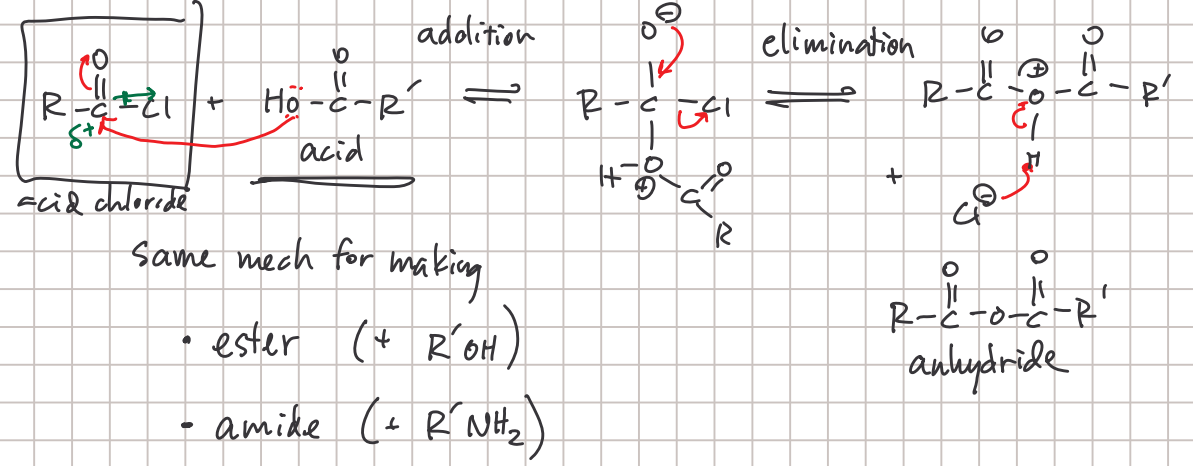
Addition / elimination mech.

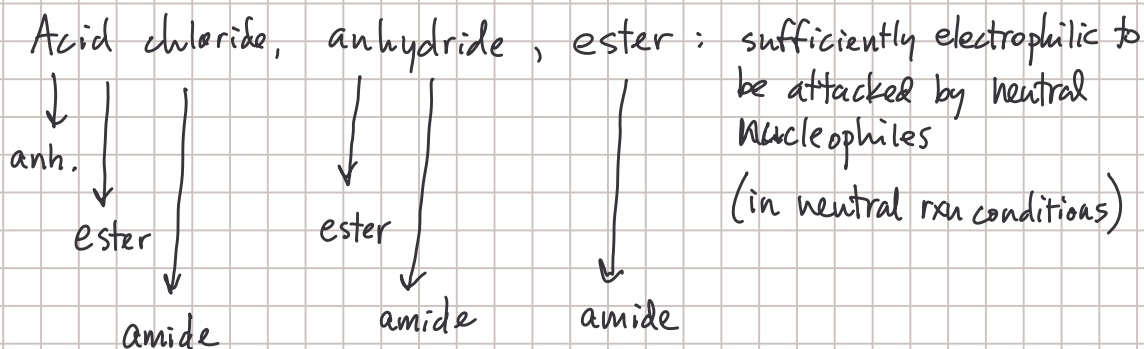


<p>most reactive</p> <p>$R-\overset{\overset{O}{\parallel}}{C}-Cl$</p> <hr/> <p>$R-\overset{\overset{O}{\parallel}}{C}-O-\overset{\overset{O}{\parallel}}{C}-R$</p> <p>$R-\overset{\overset{O}{\parallel}}{C}-OR$</p> <p>$R-\overset{\overset{O}{\parallel}}{C}-NH_2$</p> <p>$R-\overset{\overset{O}{\parallel}}{C}-O^-$</p> <p>least reactive (to addn/elim)</p>	<p>Leaving group</p> <p>Cl^- v. weak base</p> <p>$^-O-\overset{\overset{O}{\parallel}}{C}-R$</p> <p>$^-OR$</p> <p>$^-NH_2$ v. strong base</p>	<p>best leaving group</p> <p>worst leaving group</p>
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Convert a deriv into a less reactive deriv

$R-\overset{\overset{O}{\parallel}}{C}-Cl \rightarrow$ any other deriv.
acid chloride (most reactive)

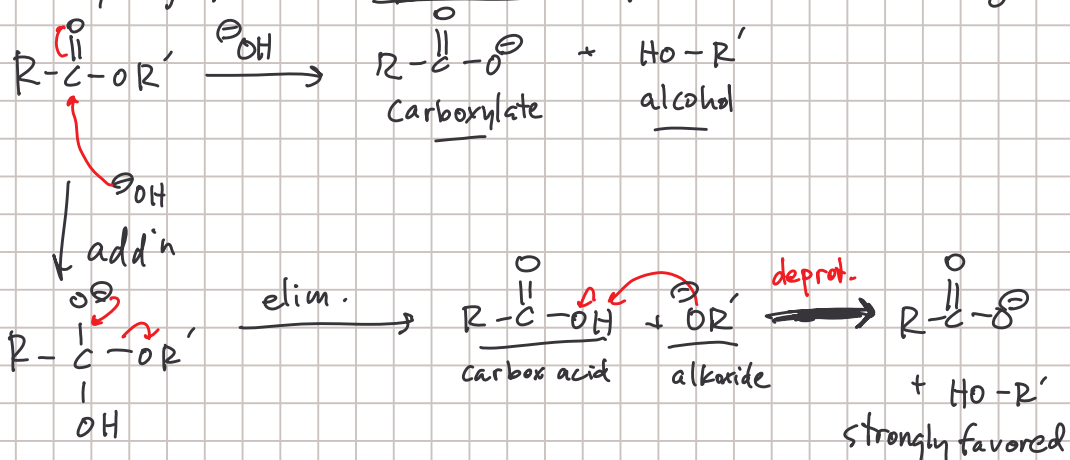




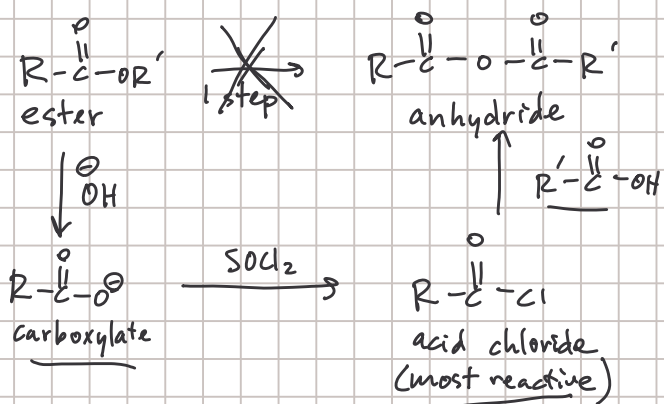
any lower-reactivity deriv must be converted to carboxylic acid first
 then converted to acid chloride by SOCl₂

Hydrolysis (addition of water)

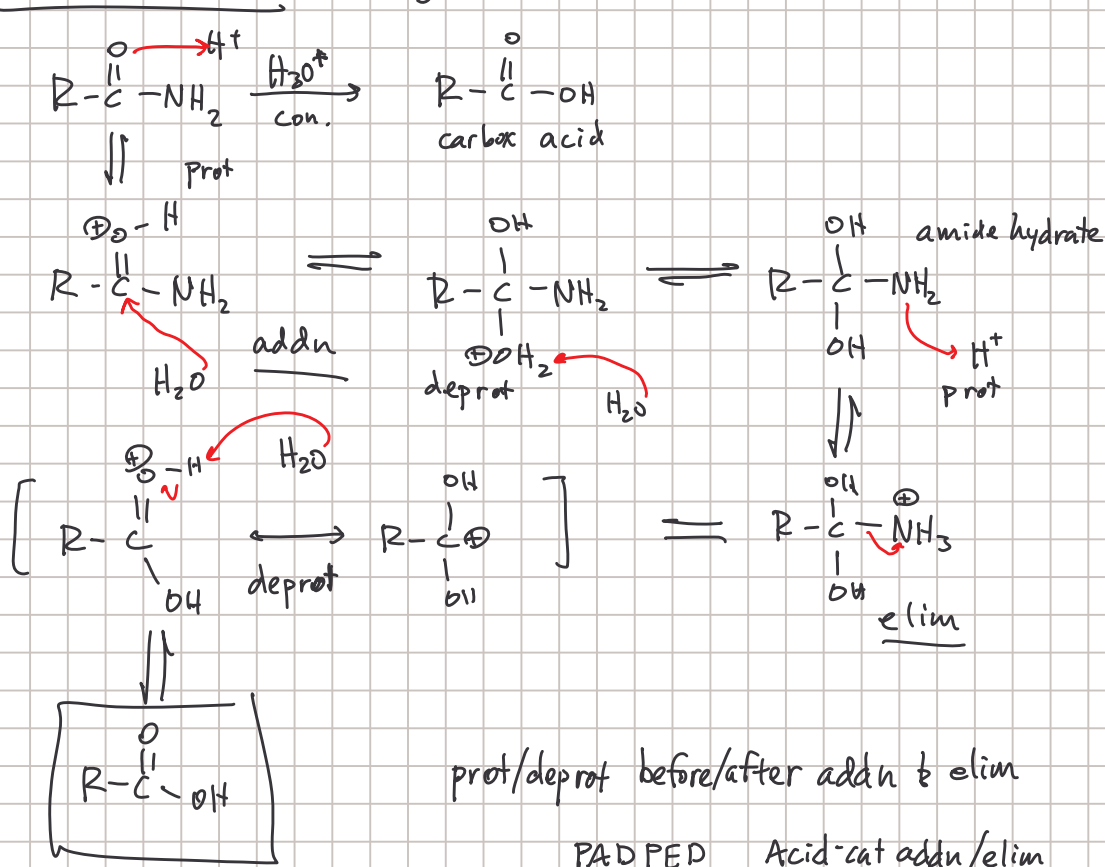
* ester hydrolysis: base-catalyzed: saponification (soap-making)



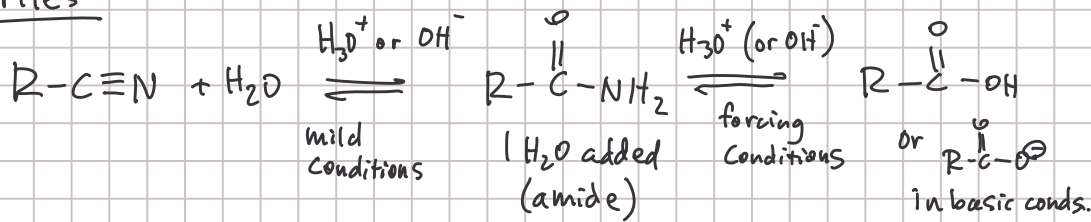
(acid-cat ester hydrolysis is reverse of Fischer esterification equilibrium-controlled, not as useful)



Amide hydrolysis strong acid conditions work best

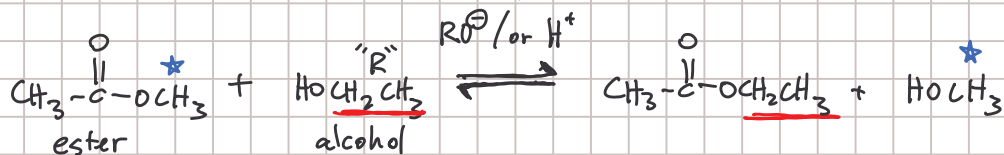


Nitriles

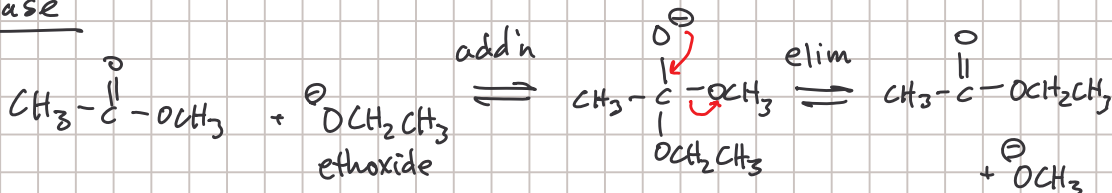


Transesterification (ester exchange)

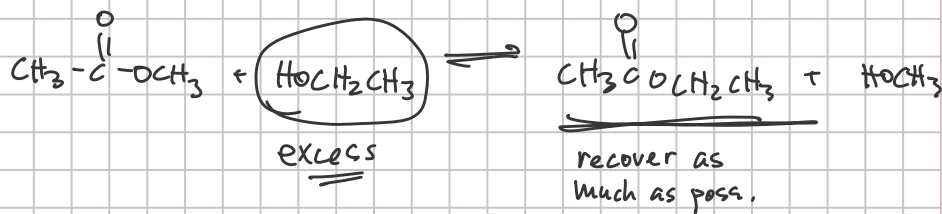
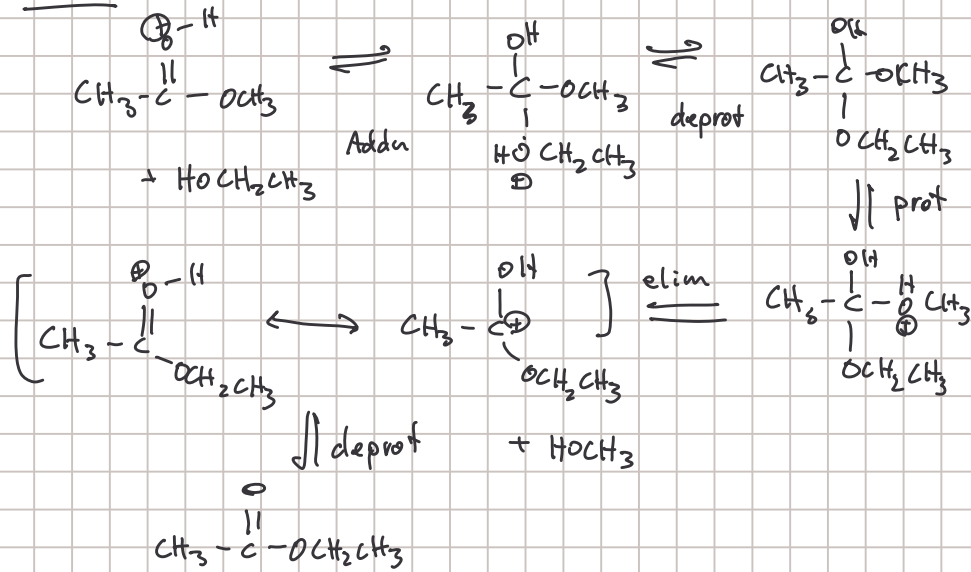
acid or base-cat - equilibrium-driven



base



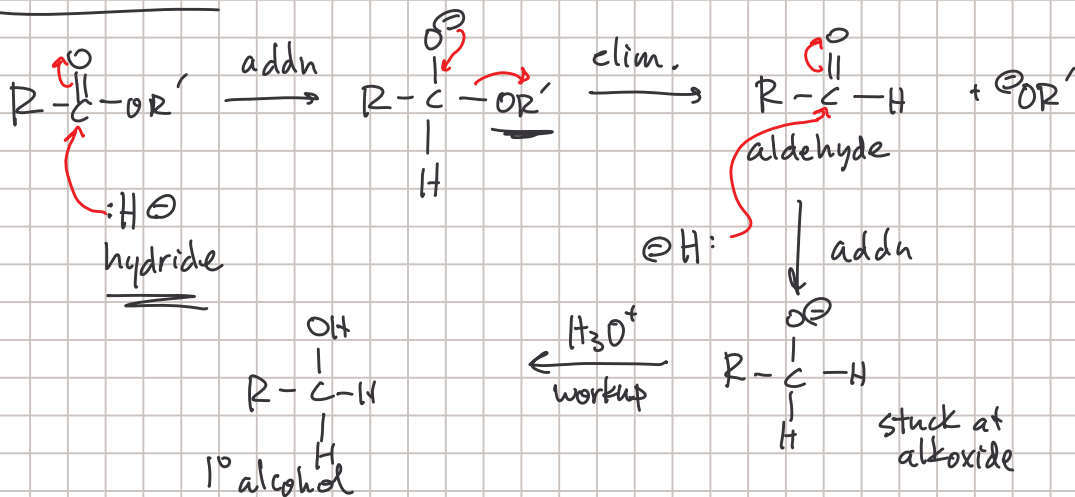
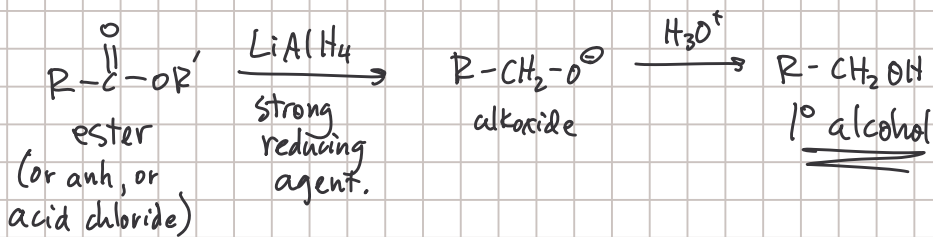
acid



Reductions

• to alcohols

ester, anh, acid chloride (good leaving groups)



reduction to amines

