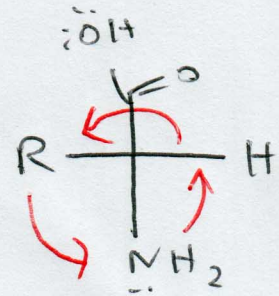
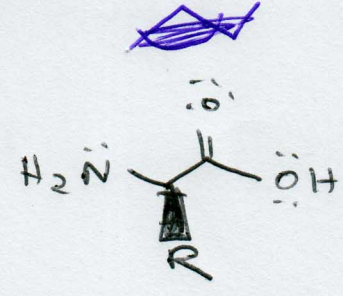
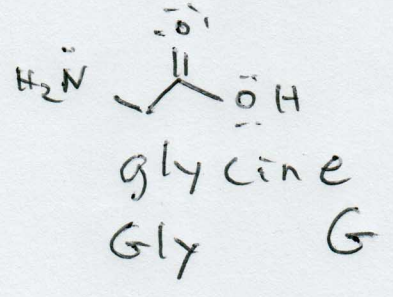
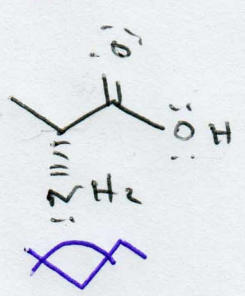
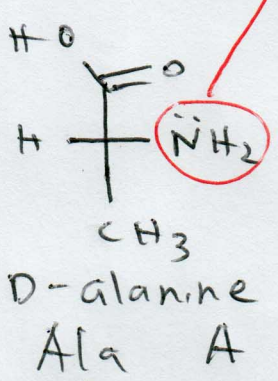


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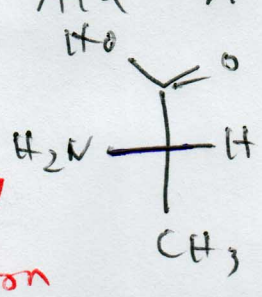
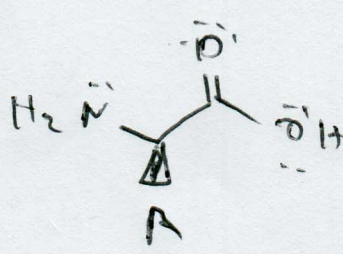
### α-amino acids



amino group on right

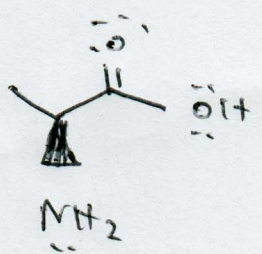


or

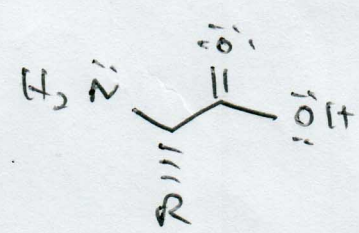


amino group on left

L-alanine

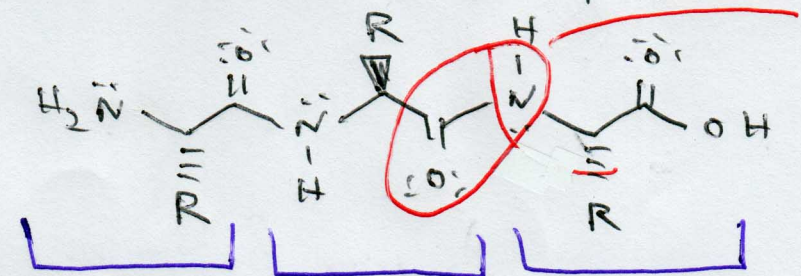


or



L-amino acids are biologically active

### peptide - amino acid polymers



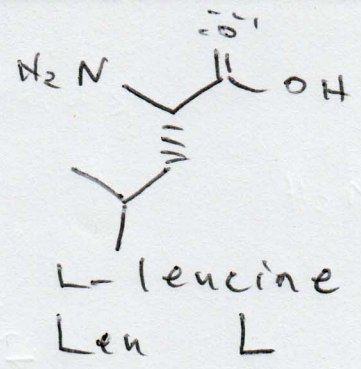
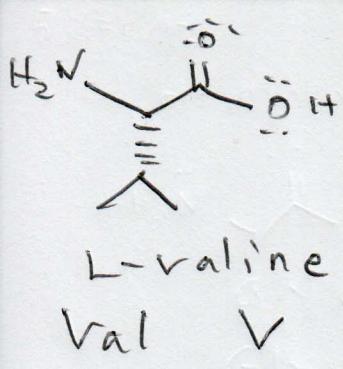
amide - relatively unreactive

3 amino acids

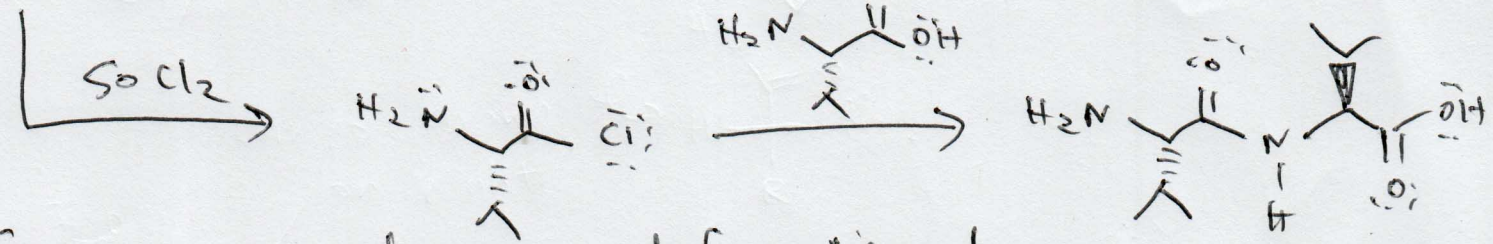
→ tripeptide

Sequence - list of amino acids in a peptide from the amino terminus to the acid terminus. (amino → acid)



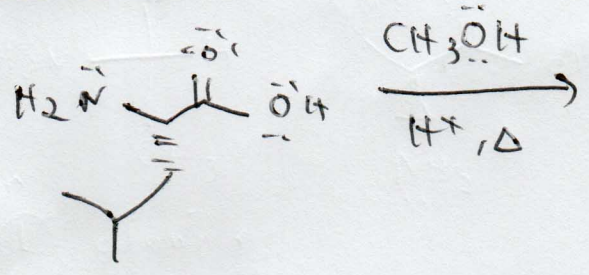


di peptide  
 $\Rightarrow$  Val-Leu

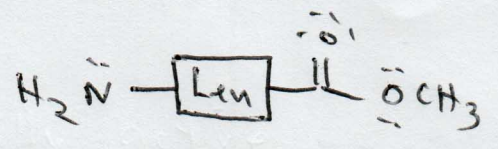


Since amino acids are difunctional, it is not possible to control the order in which amino acids add to one another without using a protecting group strategy.

C-protection



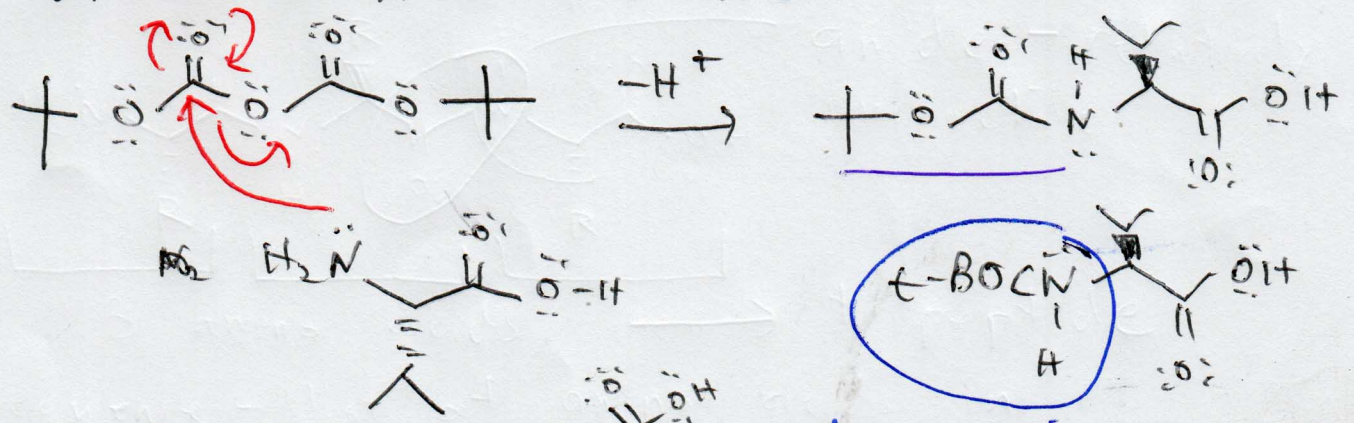
won't react w/ SOCl<sub>2</sub> or other reagents if carboxylic acid would



N-protection

di-tert-butyl dicarbonate

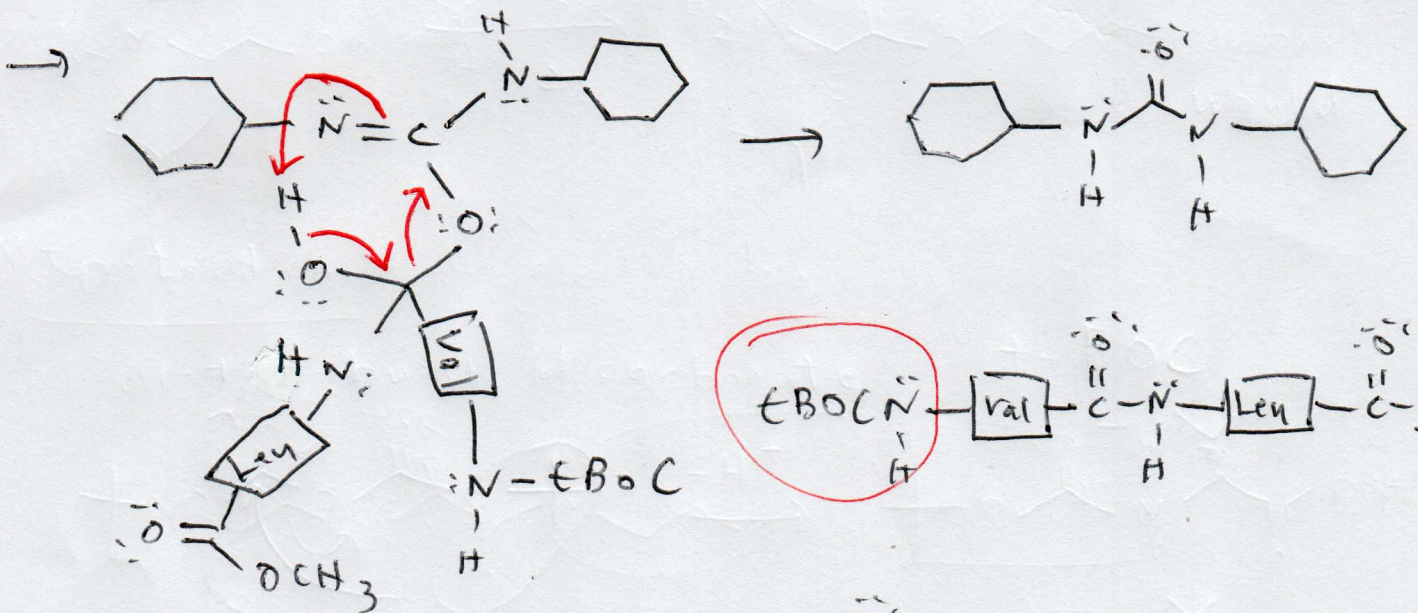
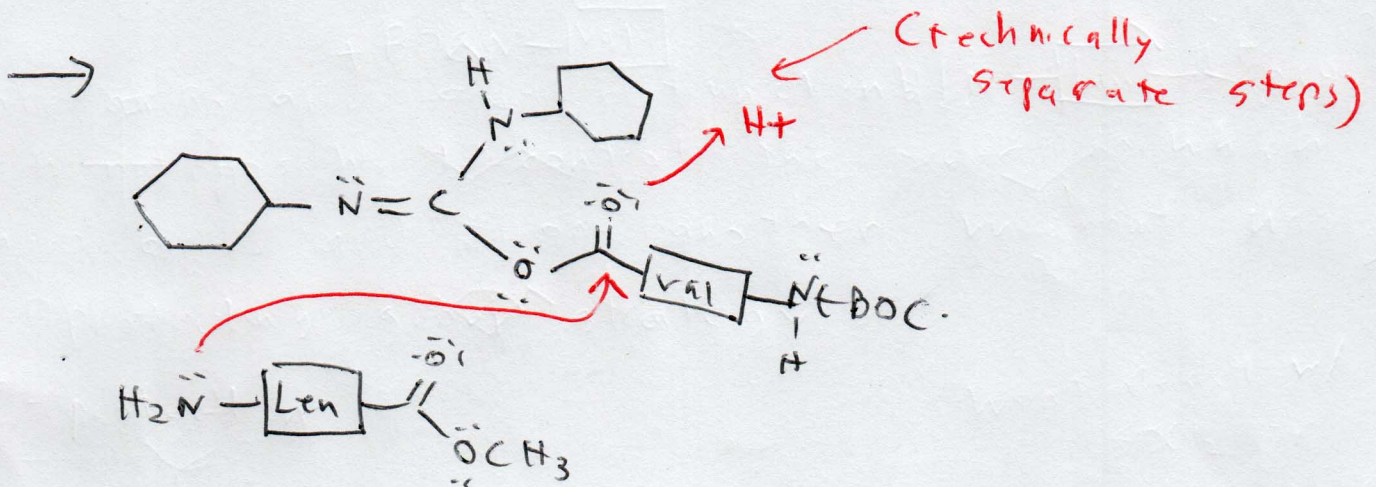
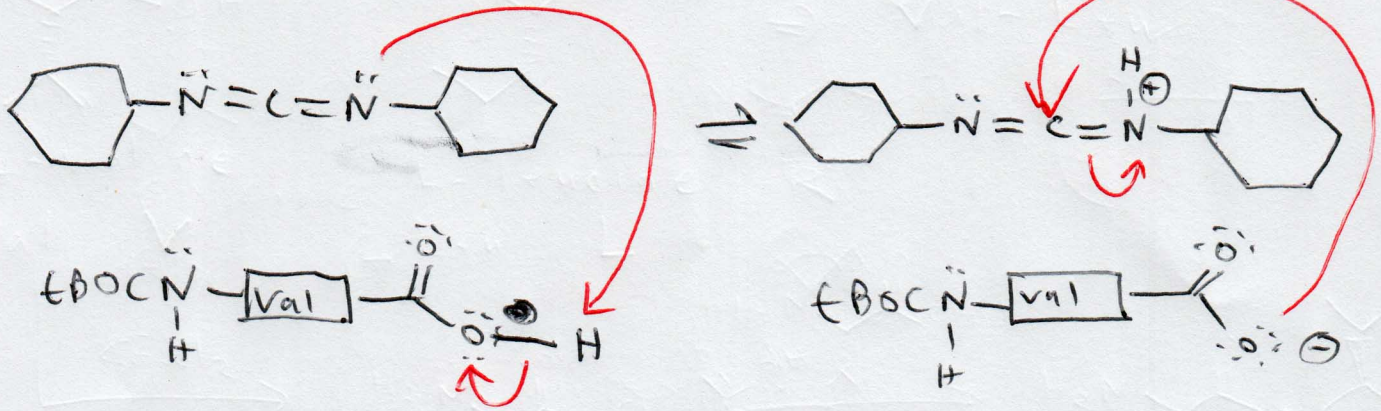
t-BOC



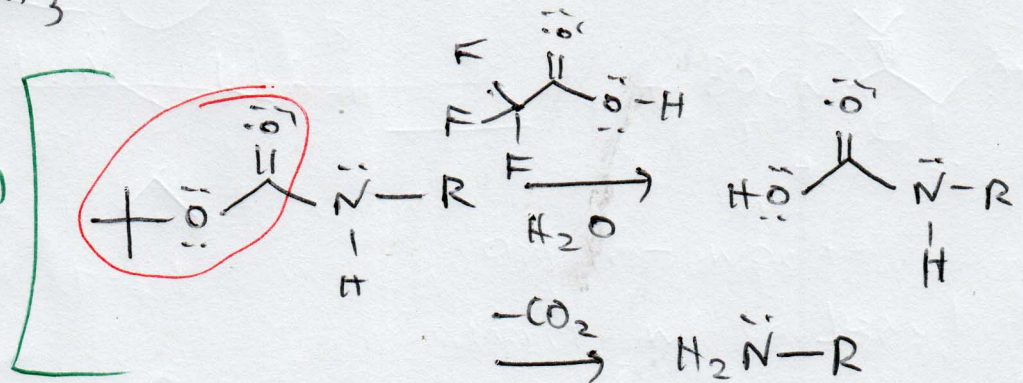
CC(C)C(NC(=O)C(C)(C)C)C(=O)O  
 amide - won't react w/ carboxylic acid



dicyclohexylcarbodiimide (DCC)

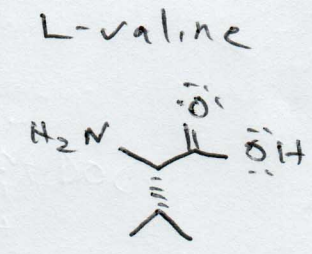
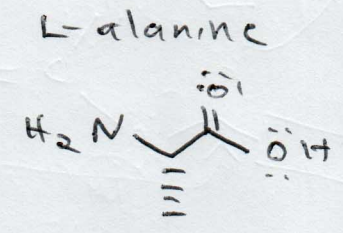
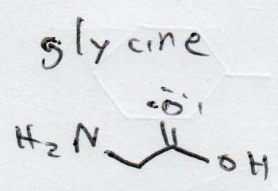
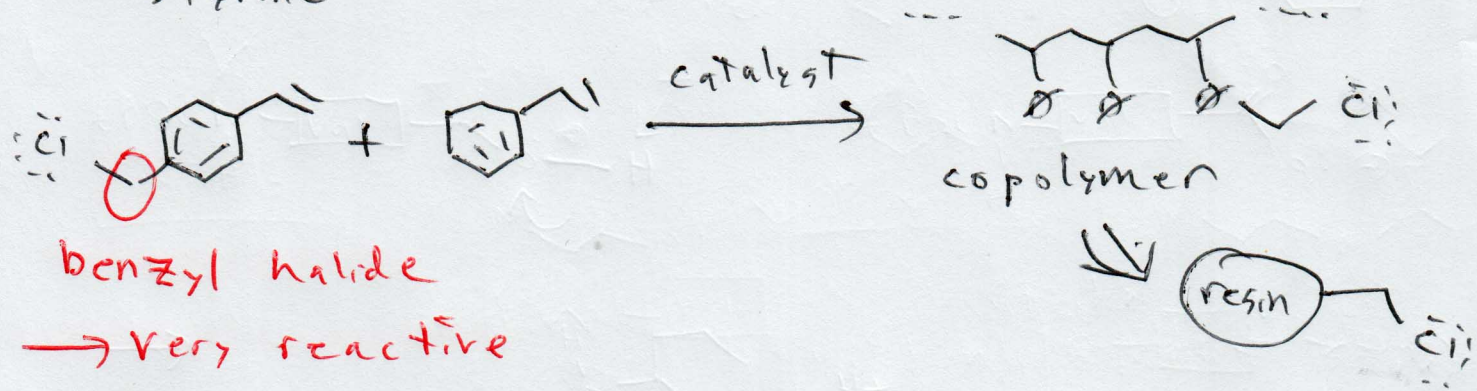
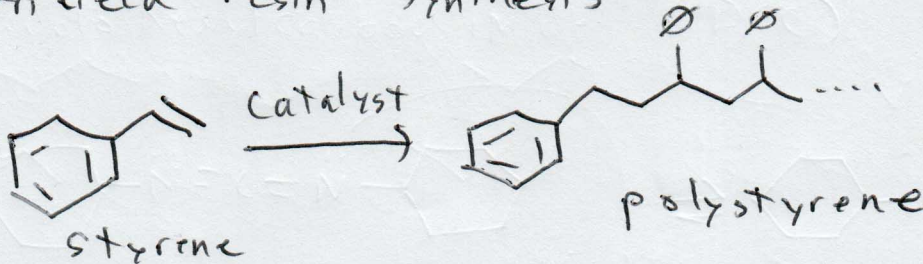


amino deprotection





Merri field resin synthesis



tripeptide VGA  $\leftarrow$  has the acid terminus that needs blocking

