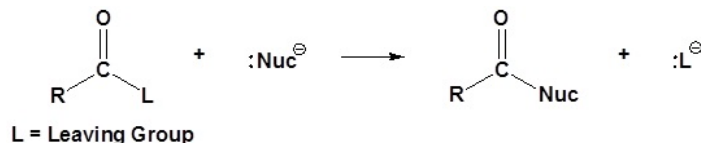


22.3: INTERCONVERSION OF ACID DERIVATIVES BY NUCLEOPHILIC ACYL SUBSTITUTION

GENERAL REACTION

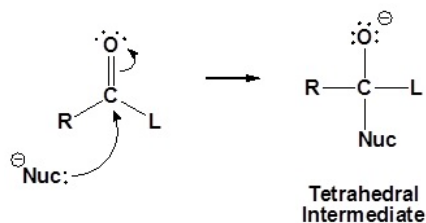
Carboxylic acid derivatives are electrophilic and can react with nucleophiles to form nucleophilic acyl substitution products. The driving force of these reactions is the stability of the leaving group shown as $:L^-$ below.



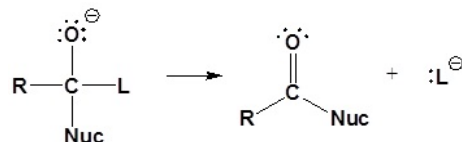
GENERAL MECHANISM

The nucleophile reacts with the electrophilic carbonyl carbon to form the tetrahedral intermediate. When the carbonyl reforms, the leaving group is lost to form the substitution product as shown in the mechanism below.

1) Nucleophilic reaction at the carbonyl




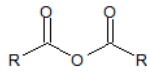


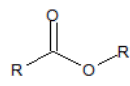
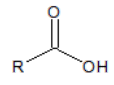
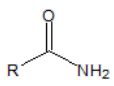
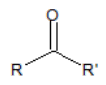
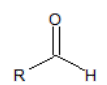
2) Carbonyl reforms and leaving group is removed



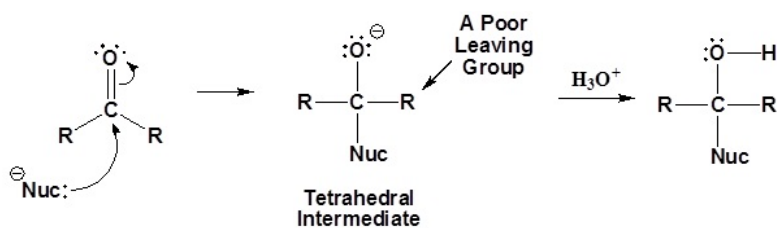
RELATIVE REACTIVITY TO NUCLEOPHILIC ACYL SUBSTITUTION

The relative reactivity of carbonyl compounds toward nucleophile substitutions is related to the stability of the leaving group - the more stable the leaving group, the more favorable the substitution reaction. Evaluating leaving group stability is analogous to evaluating conjugate base stability as shown in the table below.

Carbonyl Compounds and their Leaving Groups


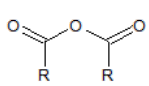
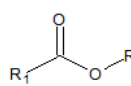
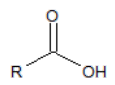
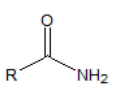
Name	Carbonyl Compound	Leaving Group	Conjugate acid of the Leaving Group	pKa
acyl chloride		Cl ⁻	HCl	-7
acid anhydride				3-5
ester		⁻ OR ₂	ROH	15-16
carboxylic acid		HO ⁻	H ₂ O	15.7
amide		⁻ NH ₂	NH ₃	36
ketone		R ⁻	RH	50
aldehyde		H ⁻	H ₂	very large

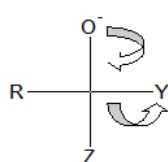
Although aldehydes and ketones also contain carbonyls, their chemistry is distinctly different because they do not contain suitable leaving groups. Once a tetrahedral intermediate is formed, aldehydes and ketones cannot reform their carbonyls because the carbide (RC⁻) and hydride (H⁻) leaving groups are too unstable. Therefore, aldehydes and ketones typically undergo nucleophilic additions and not substitutions.



Integrating all of this information into the single table below summarizes the relative reactivity of carboxylic acids and their derivatives.

Carboxylic Acid Derivatives and their Relative Reactivity

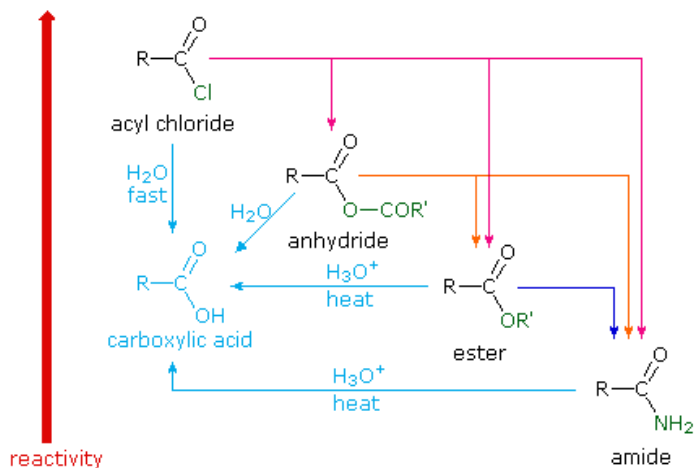
Name	Carboxylic Acid Derivative	LG	Relative Basicity	Relative Reactivity
acyl chloride		Cl^-	weakest base	most reactive
anhydride		^-O-C(=O)R		
ester		^-OR_2		
carboxylic acid		^-OH		
amide		^-NH_2	strongest base	least reactive



The weaker the base, the easier it is to expel.

ACID DERIVATIVE INTERCONVERSION

From this understanding, multiple step synthesis strategies can be developed. The derivatives with the most stable leaving groups can be used to synthesize the derivatives with the least stable leaving groups as illustrated in the diagram below.



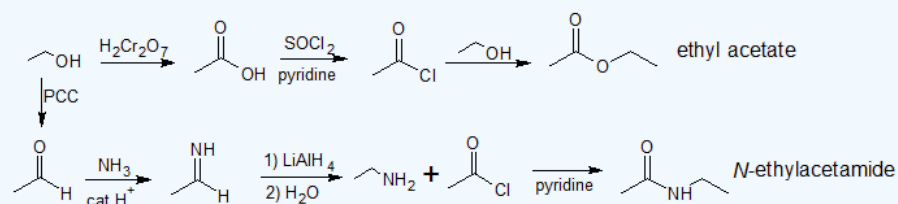
While it may appear from diagram above that the acyl chlorides are the "source" of the acid derivatives, acyl chlorides are so highly reactive that it is common to convert the carboxylic acid to the acid chloride and then immediately form the derivative. From a laboratory synthesis perspective, the reaction sequence begins with the carboxylic acid.

Exercise

3. Using ethanol as the only source of carbons in the final products, show how to synthesize ethyl acetate and N-ethylacetamide.

Answer

3.



CONTRIBUTORS AND ATTRIBUTIONS

- Prof. Steven Farmer ([Sonoma State University](#))
- William Reusch, Professor Emeritus ([Michigan State U.](#)), [Virtual Textbook of Organic Chemistry](#)

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