

23.10: CONJUGATE CARBONYL ADDITIONS - THE MICHAEL REACTION

OBJECTIVES

After completing this section, you should be able to

1. write an equation to illustrate the Michael reaction.
2. write a detailed mechanism for a given typical Michael reaction.
3. identify the product formed in a given Michael reaction.
4. identify the reagents necessary to synthesize a given compound by a Michael reaction.

KEY TERMS

- Michael reaction

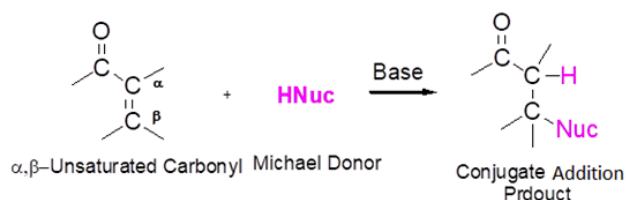
STUDY NOTES

Before studying this section in detail, you should review conjugate addition to α,β -unsaturated carbonyl compounds (Section 19.13).

Notice the wide variety of compounds that can take part in a Michael reaction. As usual, you should not feel overwhelmed by the number of different compounds that can be used, just keep in mind that in all cases the mechanism is essentially the same.

MICHAEL REACTION

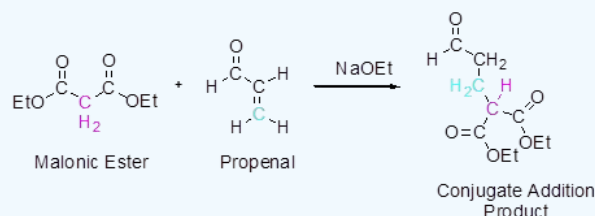
Certain nucleophiles undergo conjugate addition with the alkene of an α,β -unsaturated carbonyl compounds rather than undergo direct nucleophilic addition with the carbonyl. During conjugate addition, a nucleophile adds to the electrophilic β -alkene carbon to form a C-Nuc bond. If the starting materials contain an ester the corresponding alkoxide is used as the base in the reaction. Otherwise a hydroxide base, such as sodium or potassium hydroxide, is commonly used.



General Reaction of the Michael Reaction

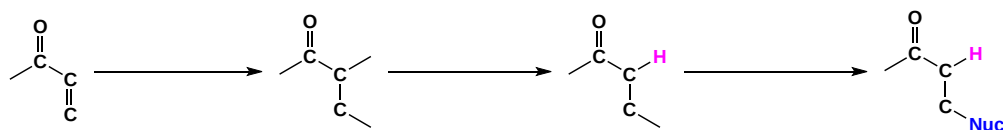
When enolate nucleophiles undergo conjugate addition with an α,β -unsaturated carbonyl the process is called a Michael Reaction. The Michael reaction works best with particularly acidic enolate donors such as malonic esters, β -keto esters, etc. Enolates which are weaker acids tend to undergo nucleophilic addition to the carbonyl rather than conjugate addition. For example, malonic ester's acidic α -hydrogens can be easily deprotonated by sodium ethoxide to form an enolate. The enolate adds to propenal to form the conjugate addition product.

EXAMPLE 23.10.1



PREDICTING THE PRODUCT OF A MICHAEL REACTION

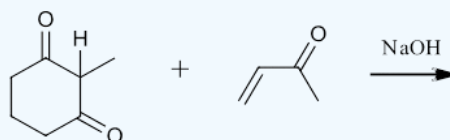
During the reaction the C=C pi bond of the α,β -unsaturated carbonyl is broken. An C-H bond is formed on the carbon in the α -position from the carbonyl. An acidic hydrogen is removed from the nucleophile and then it is used to form a C-C bond with the carbon in the β position from the carbonyl.



- In the first step, remove the double bond and add two single bonds
- Add a hydrogen to the bond adjacent to the carbonyl
- Add the nucleophile to the other end of the double bond (remember to remove a hydrogen from the nucleophile)

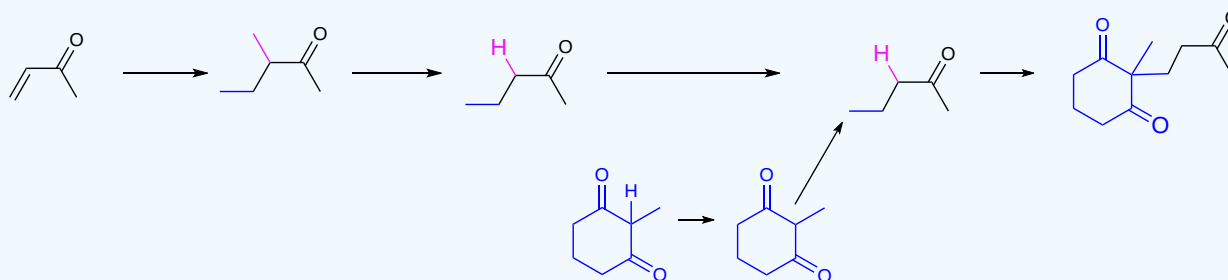
? WORKED EXAMPLE 23.10.1

What would be the product of the following conjugate addition?



Solution

- Break the double bond and add two single bonds
- Add a hydrogen to the bond adjacent to the carbonyl
- Remove a hydrogen from the nucleophile.
- Add the nucleophile to the other end of the double bond.



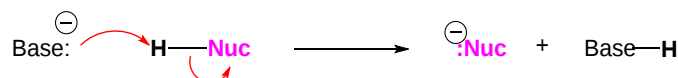
MECHANISM OF MICHAEL REACTION

During a Michael reaction the enolate acts as a nucleophilic donor and the α, β -unsaturated carbonyl acts as the electrophilic acceptor. The mechanism is a mixture of an alpha-substitution for the enolate and a conjugate addition for the α, β -unsaturated carbonyl.

Notice that the Michael Reaction does not require the deprotonation of the product to push the reaction towards completion. The reaction is thermodynamically favorable because the C-C bond formed in the product is stronger than the C=C bond in the starting material. This means that, unlike the Claisen condensation, this reaction only requires a catalytic amount of reaction base.

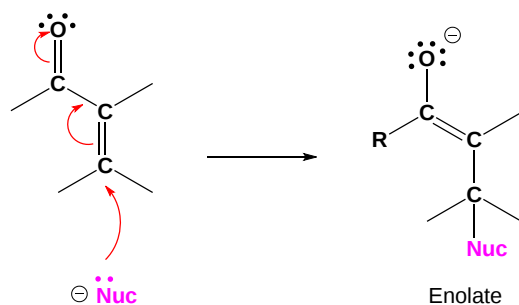
STEP 1: DEPROTONATION

The alkoxide base removes an alpha-hydrogen to form an enolate nucleophile.



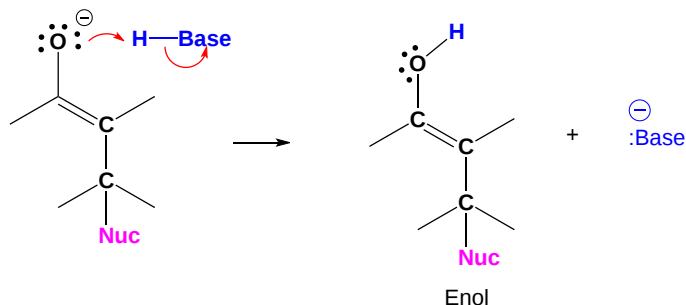
STEP 2: NUCLEOPHILIC ATTACK ON THE CARBON B TO THE CARBONYL

The enolate nucleophile then adds to the electrophilic β -carbon of the α, β -unsaturated carbonyl. Conjugation pushes the C=C pi bond electrons onto the carbonyl oxygen, forming a new enolate enolate.



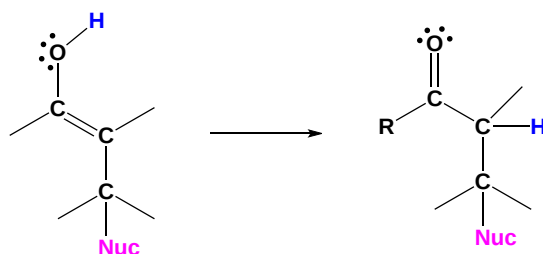
STEP 3: PROTONATION

The alkoxide is protonated, forming an enol.



STEP 4: TAUTOMERIZATION

Tautomerization converts the enol back into a carbonyl forming the neutral conjugate addition product.



The Michael reaction can be performed with a wide variety of α , β -unsaturated carbonyl electrophilic acceptors and enolate donors. Michael acceptors such as α , β -unsaturated ketones, aldehydes, esters, amides, and nitro compounds can all participate in this reaction. Michael donors with particularly acidic α -hydrogens, such as β -diketones, β -keto esters, β -keto nitriles, α -nitro ketones, malonic esters, and nitro compounds can be used.













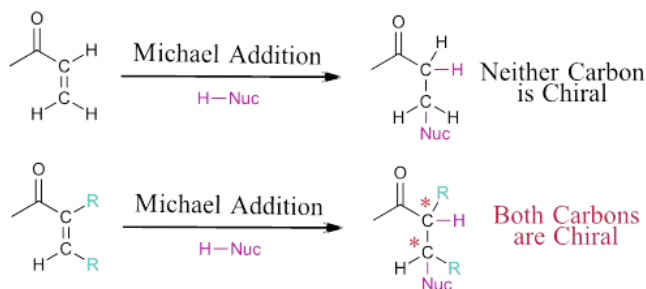
Michael Donors (aka nucleophile)	Michael Acceptors (aka electrophile)
 β -diketone	 α,β -unsaturated aldehyde
 β -keto ester	 α,β -unsaturated ketone
 β -keto nitrile	 α,β -unsaturated ester
 α -nitro ketone	 α,β -unsaturated amide
 malonic esters	 α,β -unsaturated nitrile
 nitro compound	 nitroethylene

Table 23.10.1 : Possible Donors and Acceptors Which can be used in a Michael Reaction

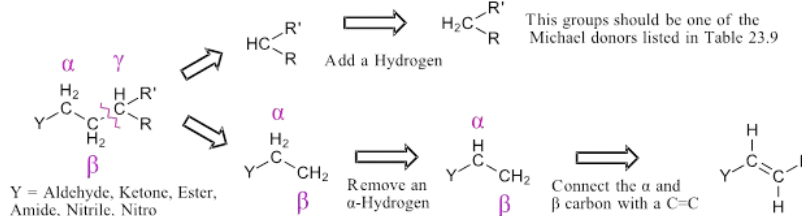
STEREOCHEMICAL CONSIDERATION OF MICHEAL REACTIONS

During a Michael reaction two sp^2 hybridized carbons are both converted to sp^3 hybridization. Both of them have the possibility of creating a chiral carbon. Michael reactions are often performed using an α, β -unsaturated carbonyl with only hydrogen substituents on the alkene to prevent the formation of chiral carbons. If either carbon in the alkene has an alkyl substituent it will likely form a chiral carbon in the Michael reaction product. The α -carbon of the enolate donor also has the possibility of forming a chiral carbon as discussed in **Section 22-6**.



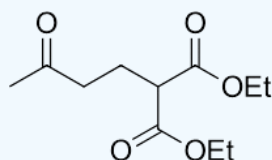
PLANNING A SYNTHESIS USING A MICHAEL REACTION

The true utility of the Michael reaction is seen when considering its use in a synthesis. The reaction can be used to prepare aldehydes, ketones, esters, amides, nitriles, and nitro compounds. A target molecule can possibly be made using a Michael reaction if it contains one of these functional groups and an alkyl chain at least three carbons long. The key bond cleavage is a C-C bond between β and γ carbons from a carbonyl-like group. The fragment with the Y group loses an α -hydrogen and then forms a C=C bond between the α and β carbon. The carbon of the other fragment gains a hydrogen. This fragment should possess an acidic α -hydrogen and should be made up of Michael donor fragments such as those listed in **Table 23.10.1**. In some cases, there may be more than one fragmentation variation. Each variation should be investigated and the possible starting materials compared for viability.



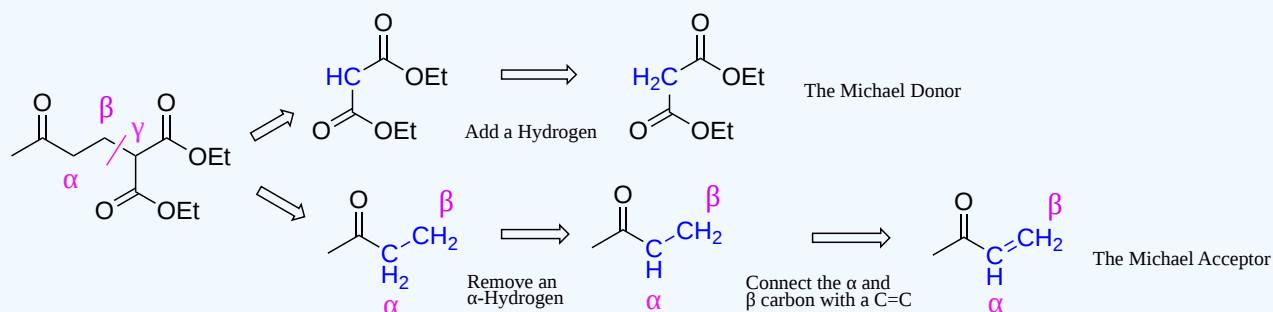
? EXAMPLE

How could the following molecule be synthesized using a Michael Reaction?

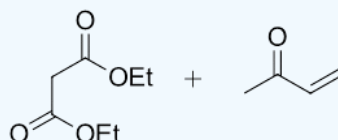


Answer

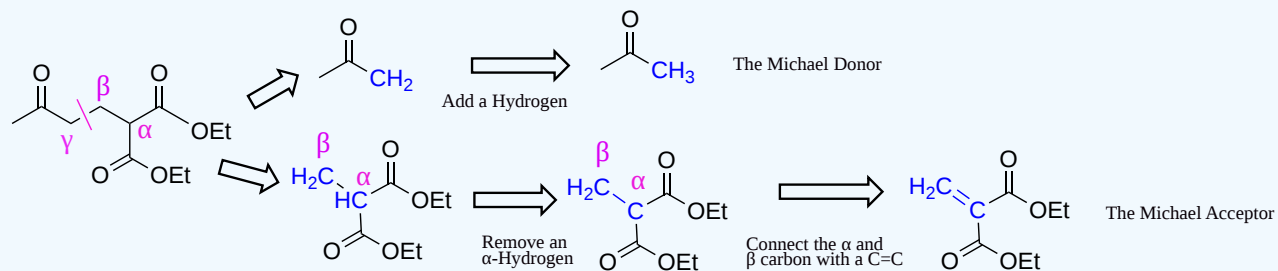
Pathway 1



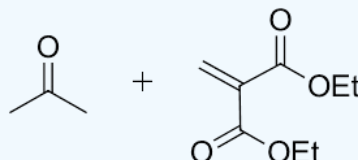
Starting Materials 1



Pathway 2

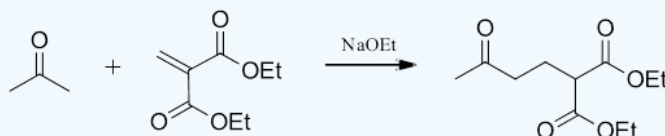


Starting Material 2

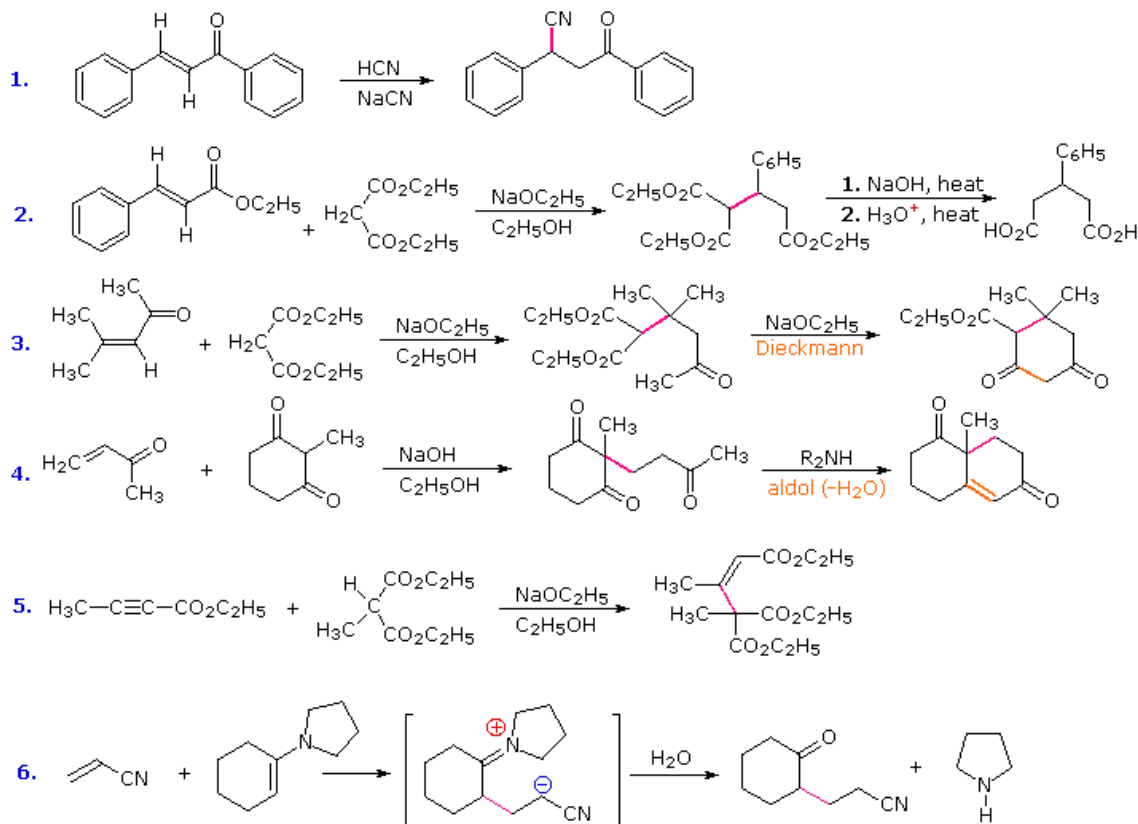


Solution

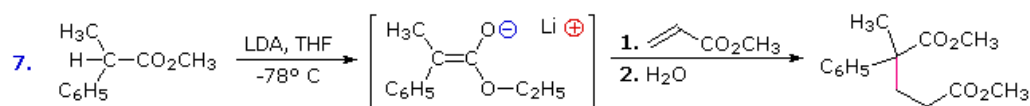
When comparing the two sets of possible starting materials of pathway 1 are preferred. The Michael enolate donor is particularly acidic because its conjugate base is stabilized by two carbonyls. The Michael enolate donor for pathway 2 does not have the same acidity. Consequently, the enolate would preferably undergo nucleophilic addition rather than conjugate addition. This makes the starting materials of pathway 1 preferred for the synthesis of the starting material.



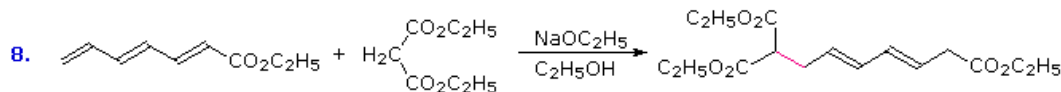
In combination with alkylations and condensations, the Michael reaction may be used to construct a wide variety of complex molecules from relatively simple starting materials. The carbon nucleophiles used in the following examples include cyanide ion, sodium diethylmalonate and the conjugate base of cyclohexane-1,3-dione. These anions are sufficiently stable that their addition reactions may be presumed reversible. If this is so, the thermodynamic argument used for hetero-nucleophile additions would apply here as well, and would indicate preferential formation of 1,4-addition products. Cyanide addition does not always follow this rule, and aldehydes often give 1,2-products (cyanohydrins). In each case the initial reaction is a Michael addition, and the new carbon-carbon bond is colored magenta. Any subsequent bonds that are formed by other reactions are colored orange.



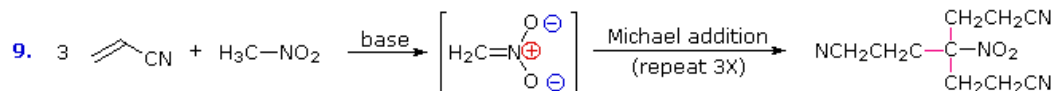
Enamine nucleophiles are good Michael donors. Hydrolysis is necessary to obtain the ketone product.



Weak carbon acids may be converted to their enolate anions prior to addition of the Michael acceptor.



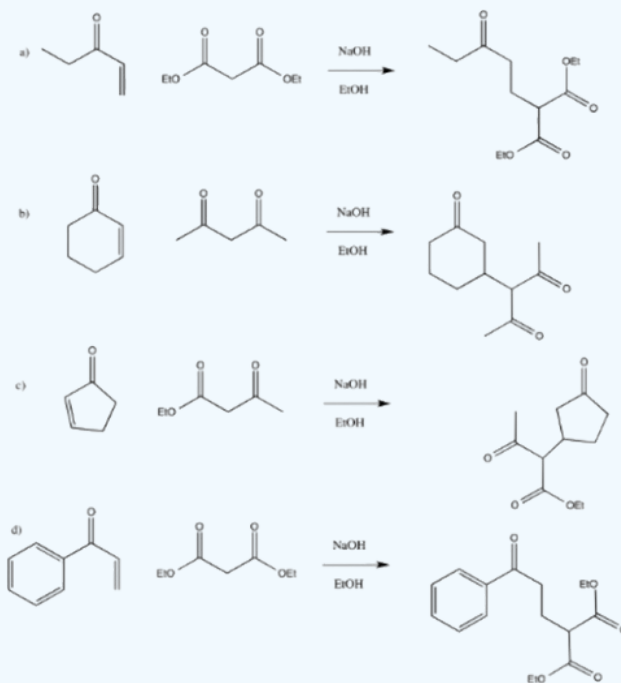
Extended vinylous Michael acceptors have been used. This is a 1,6-addition (1,8 if the enol is counted).



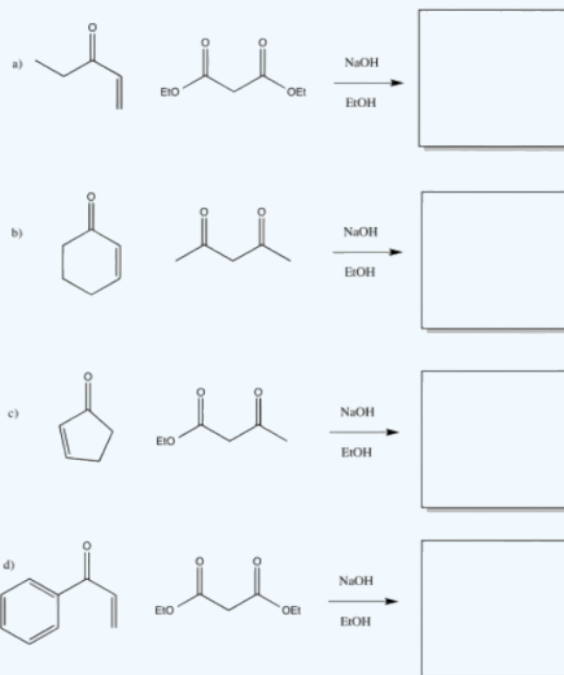
Nitroalkanes have acidic α -hydrogens and make excellent Michael donors.

? EXERCISE 23.10.1

Provide the products of the following Michael additions:

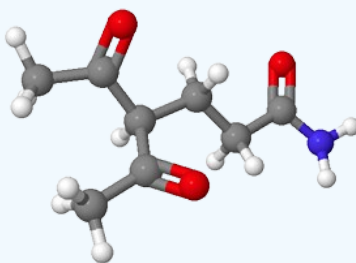


Answer

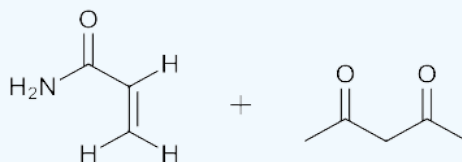


? EXERCISE 23.10.2

What would be the starting materials required to make the following molecule using a Michael reaction? Red = Oxygen, Grey = Carbon, White = Hydrogen, Blue = Nitrogen.



Answers



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