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Brain cancer patient wins Gusher Marathon

By **Avi Zaleon**

Updated March 11, 2013 11:27 a.m.



Iram Leon and daughter Kiana Leon finish the Gusher Marathon in 3 hours 7 minutes and 35 seconds taking first place in the full marathon on Saturday, March 9, 2013. Photo taken: Randy Edwards/The Enterprise

A man racing against time crossed the finish line. His daughter, snugly sitting in a stroller he had been pushing for just over 26 miles, gained a memory that may live longer than her father.

"This is supposed to eat away at my memory in the end," Iram Leon said of the cancer in his left temporal lobe. "But I hope this memory is one of the last things to go and one she never loses."

Leon is 32 years old. He said his doctors have told him, "we're probably not going to beat this. We're just hoping to get you to 40."



Hello Dr. Iverson,

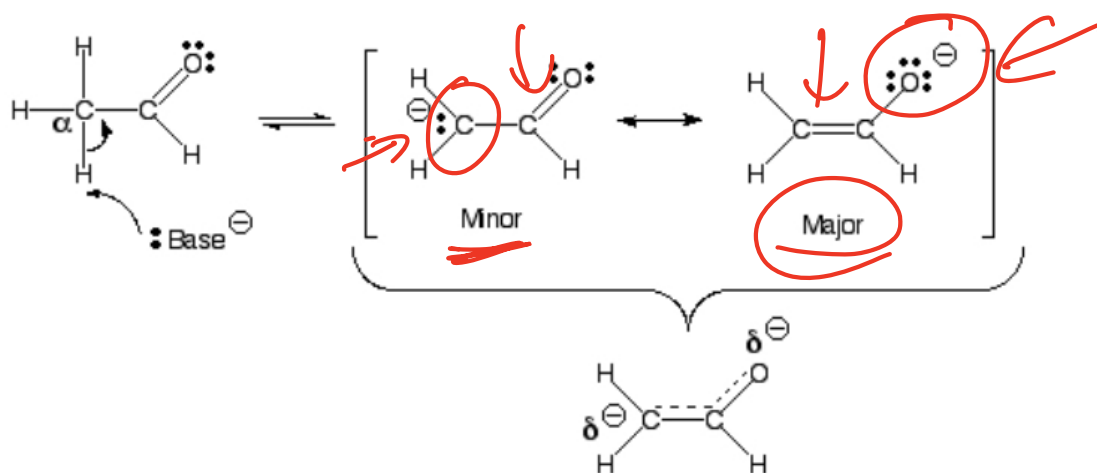
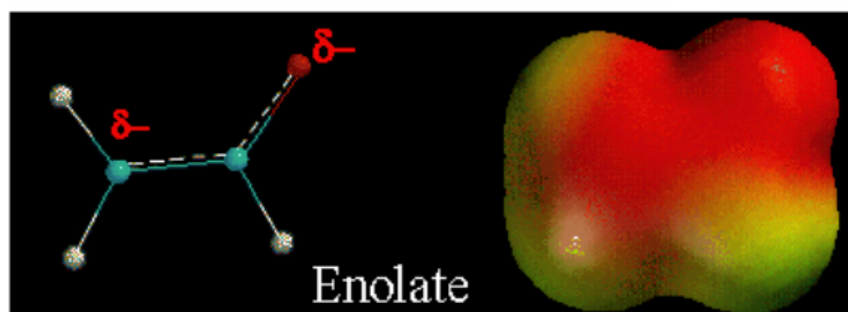
You may not remember me, but I was in your organic chemistry class last semester.

This past summer I was diagnosed with lymphoma cancer. Initially I had lost all hope, I kept asking myself "why me?" and kept thinking of all things I hadn't accomplished in my lifetime. Nevertheless, I soon got over that fact and started my chemotherapy treatments. Each treatment got worse and worse as I experienced more and more of the side effects. At night I couldn't fall asleep from all psychological and financial stress, couldn't eat because of mouth sores, and when I did eat I would feel sick and nauseated. It wasn't until my third treatment that I remembered the many times you told the class that running could help quality of life. It took a couple of weeks for me to convince myself to start running but I eventually started slowly. I never thought how great of an affect physical activity could have. I was never obese so I never gave running or cardio any thought. As I started running on a regular basis I started seeing my symptoms disappear slowly. Soon when I would come back from running I suddenly had an appetite, regardless of the mouth sores I was hungry enough to eat. My sleeping schedule was started falling into place because I was so tired after running. My stress levels decreased enough that I could see the difference. Best of all it gave me something to do during my days at home, saving me from depression.

Running saved my life Dr. Iverson.

Thanks again,

Enolates as nucleophiles

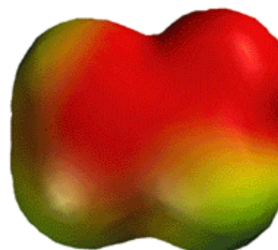
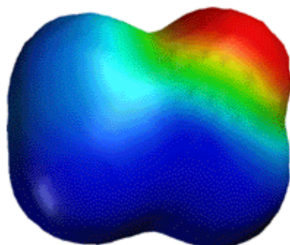
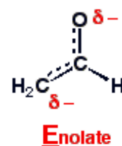
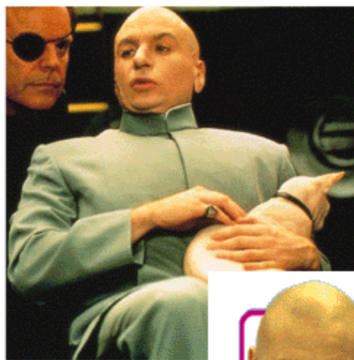
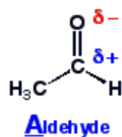


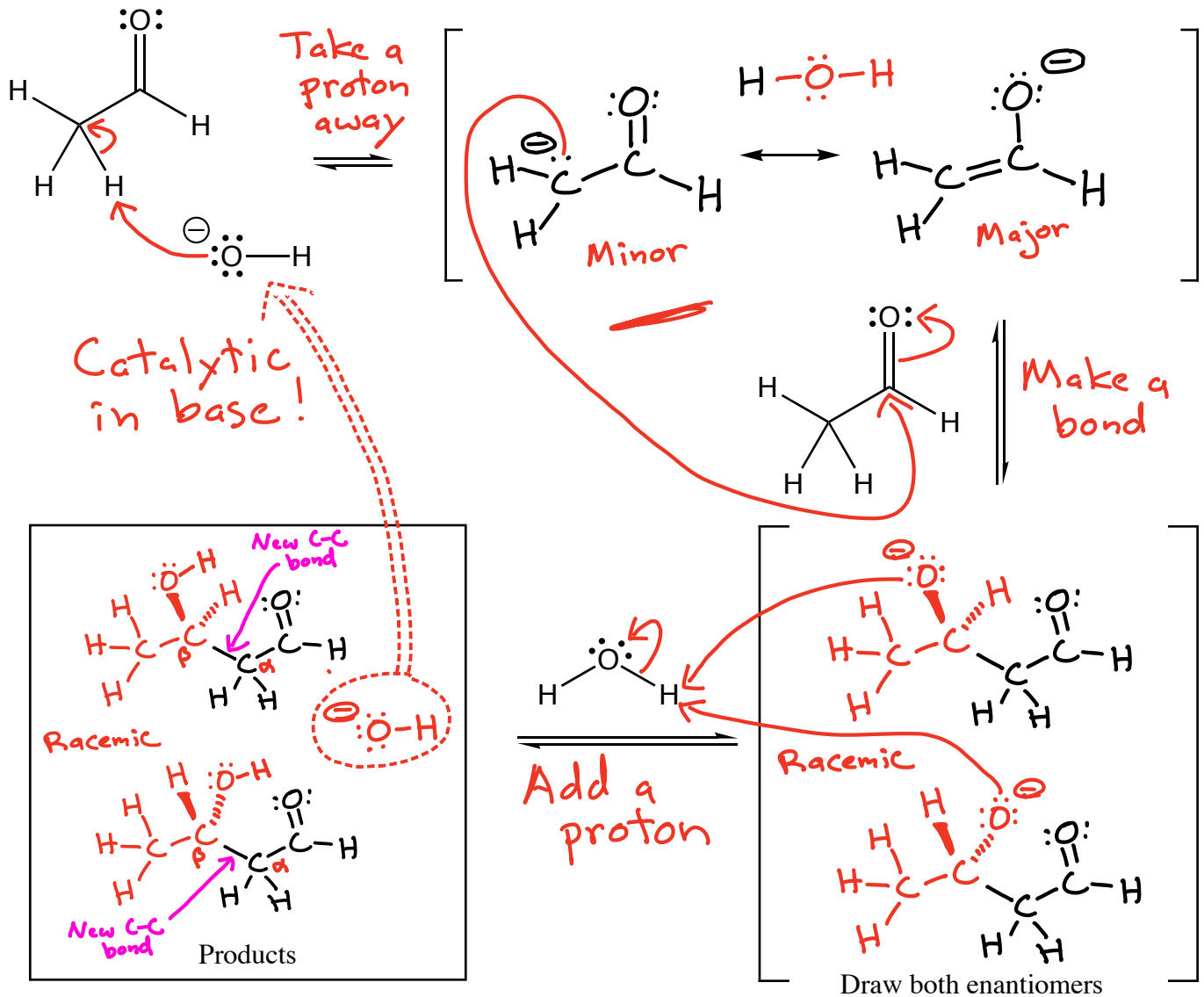
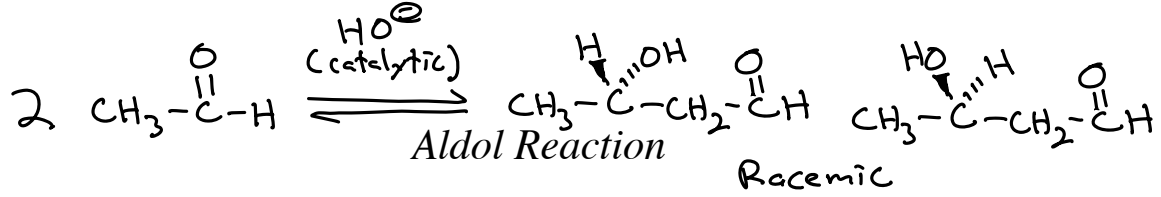
A) Enolates are resonance stabilized, with a partial negative charge on carbon and oxygen.

B) Enolates are nucleophiles, so they could react at either the carbon atom or oxygen atom. The partial negative charges give them the **opportunity** to react at either the carbon or oxygen.

C) Reaction at the carbon atom gives the final product a C=O bond, while reaction at the oxygen atom gives the final product a C=C bond. However, C=O bonds are stronger than C=C bonds, **so the motive is to react at the carbon atom with most electrophiles.**

Once Again, A Movie Ripping Off Chemistry

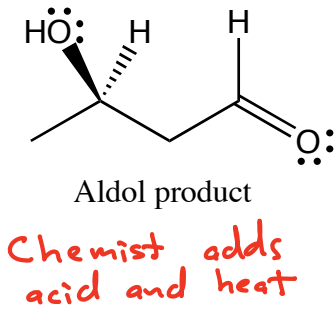




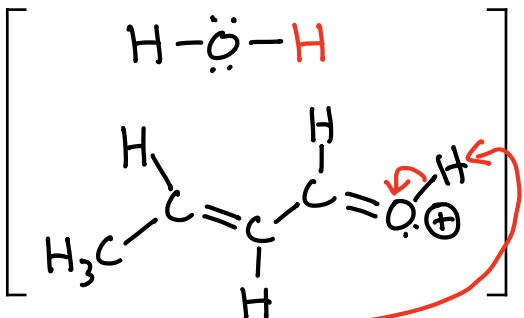
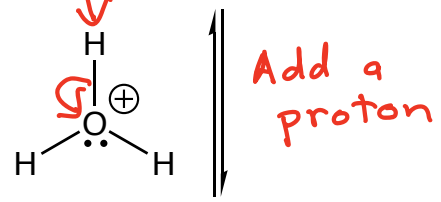
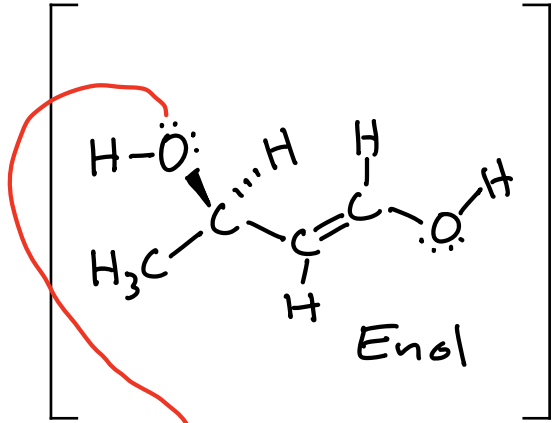
KRE → β-hydroxy aldehyde
 with a new C-C
 bond between the
 aldehyde α and β
 carbons

Mechanism
 A

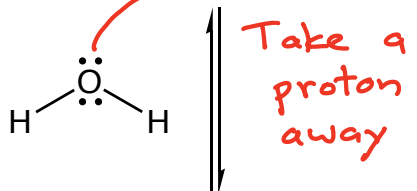
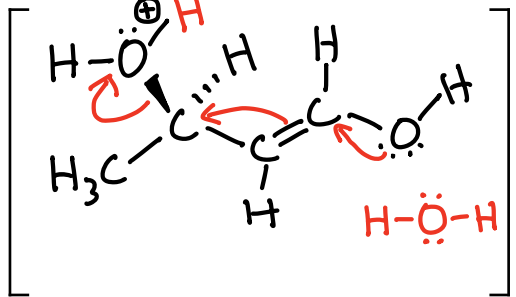
Acid catalyzed dehydration



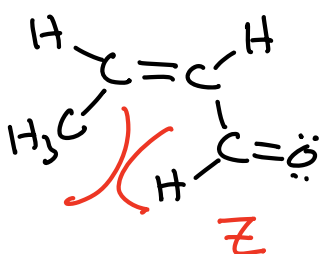
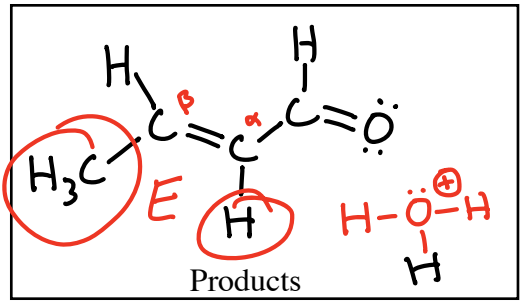
tautomerization



Break a bond



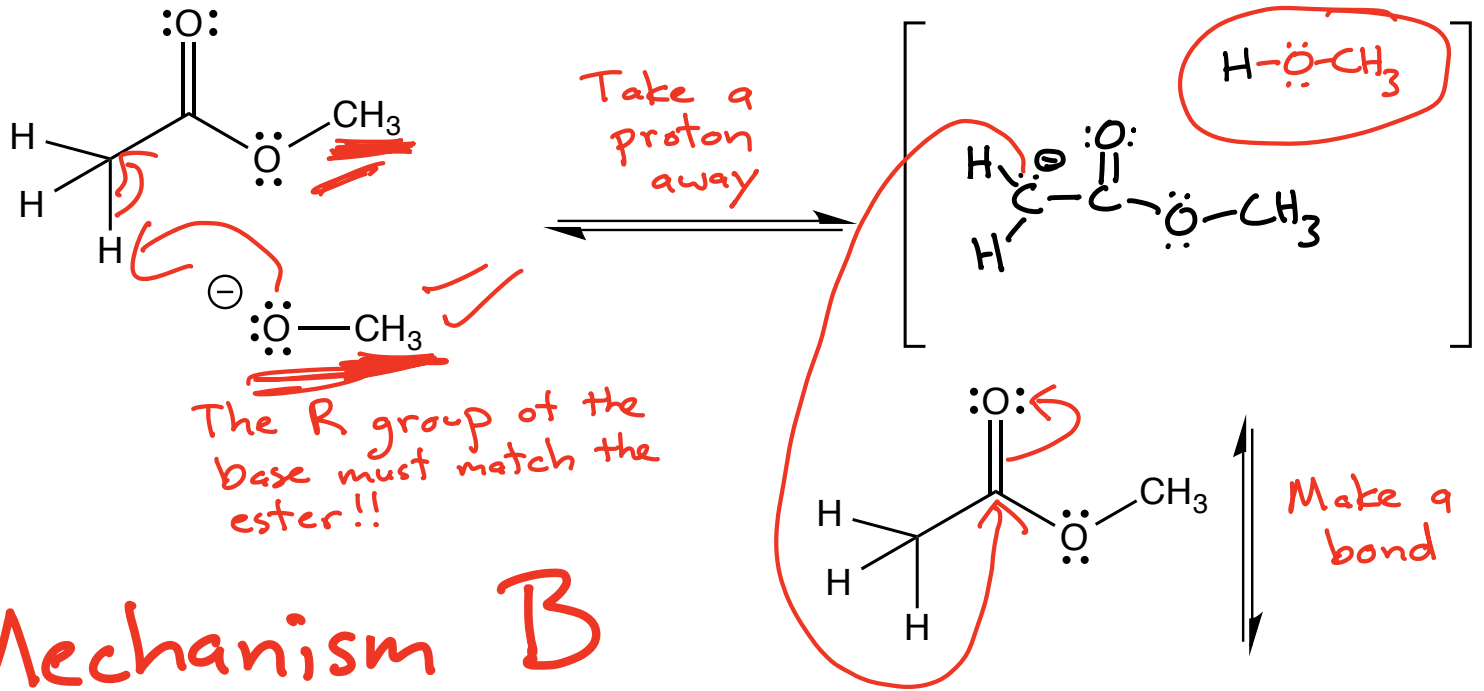
KRE → α,β-unsaturated aldehyde → the C=C is where the new C-C bond is located



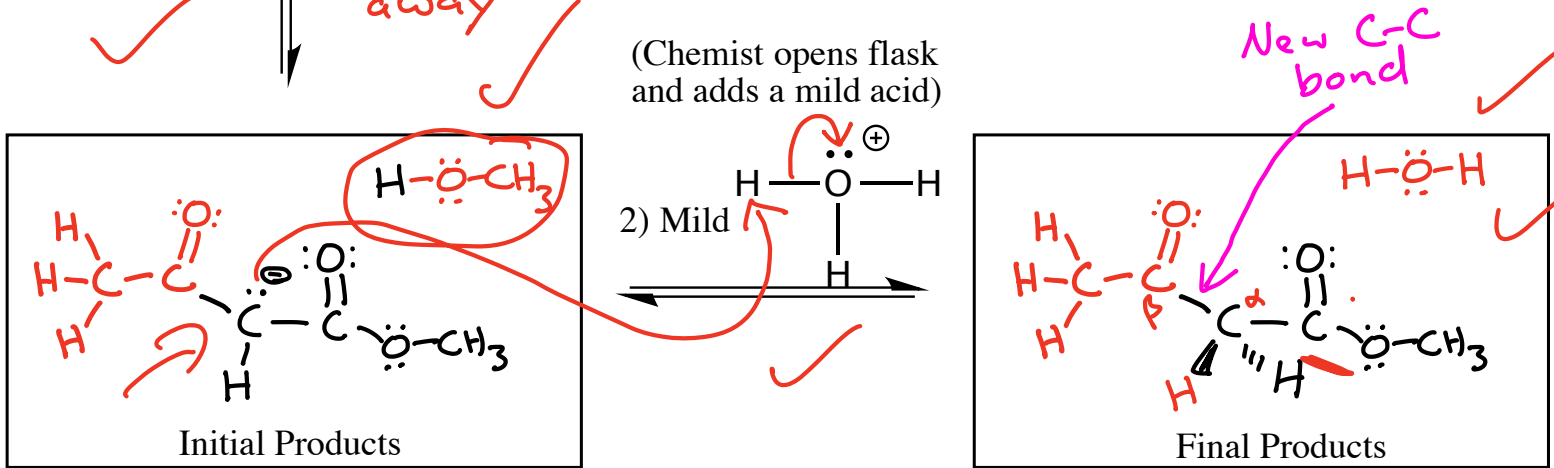
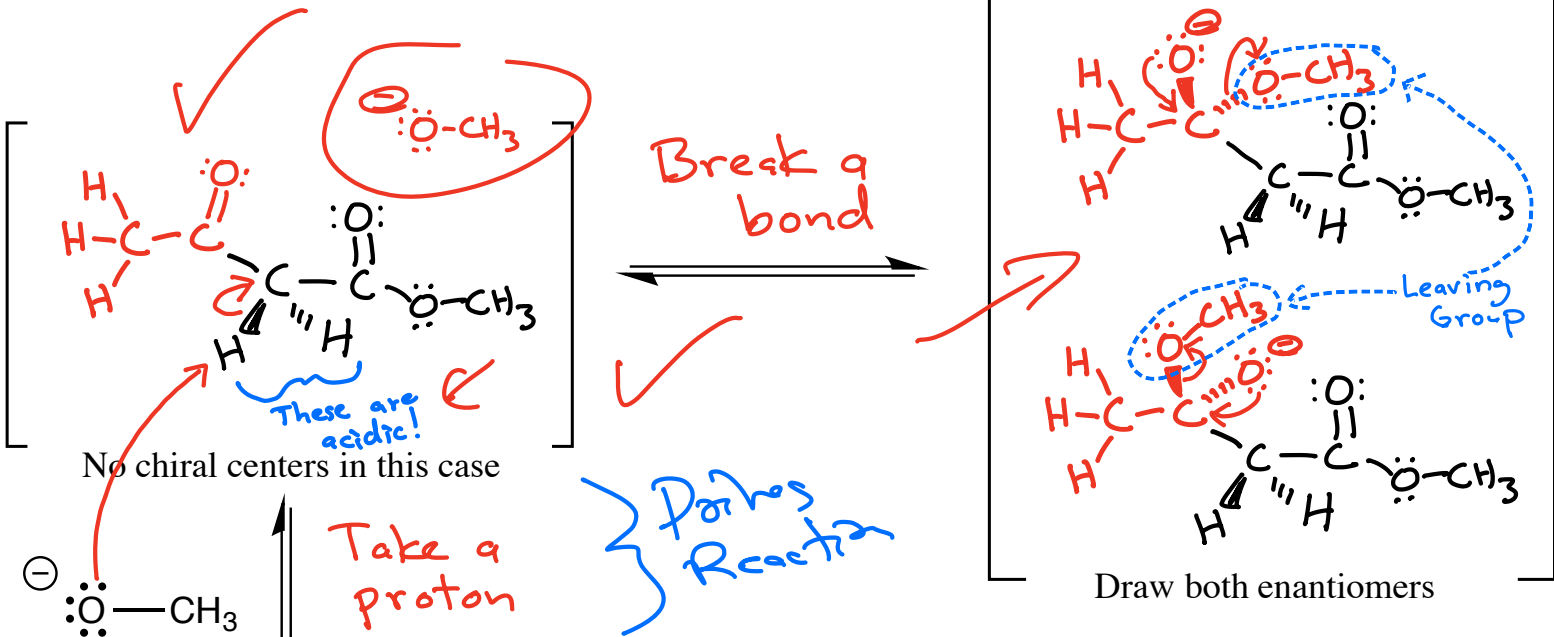
THIS IS UNIQUE TO THIS EXAMPLE
 ↓
 USUALLY BOTH E AND Z ARE FORMED

Not much of the Z product is formed because it has significantly more steric strain than E

Claisen Condensation → "Aldol with Esters"



Mechanism B

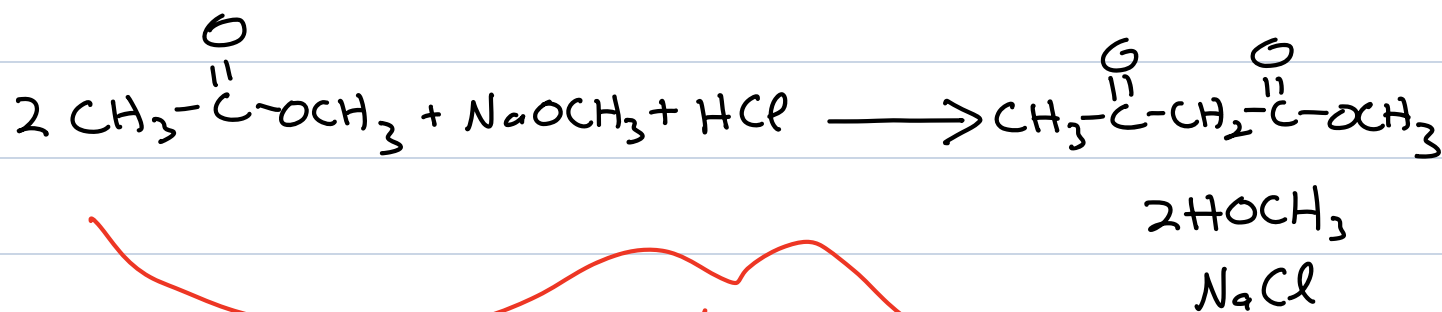


This is a much more stable anion compared to $\ominus\text{OCH}_3$, providing a strong driving force (motive) for the Claisen condensation reaction

- KRE \rightarrow A β -keto ester with a new C-C bond between the α and β carbons

Before we add acid \rightarrow the last step drives the reaction because we make a relatively stable anion.

Balanced Equation for the Overall Process

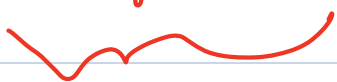
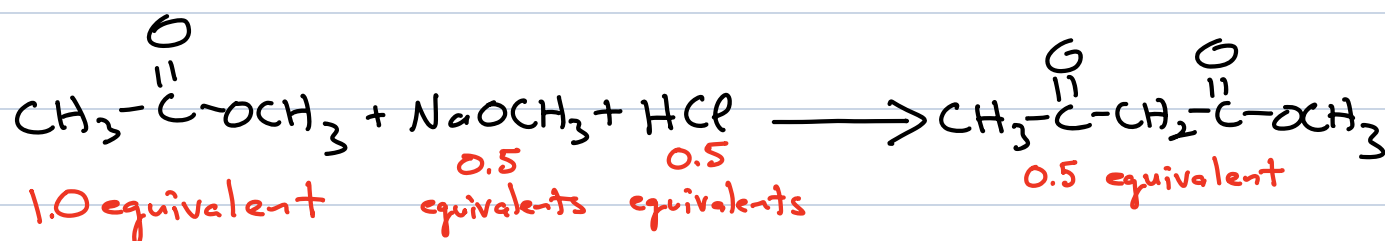


This is the balanced equation that is explained by the mechanism

With the balanced equation in hand we can set up a reaction properly in the lab because we know how much of each reactant is needed → For this we use the notation of "equivalents"



Notation we use for the lab and therefore synthesis and "box" problems on the exams

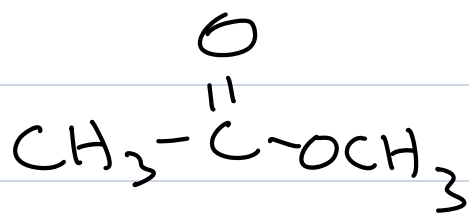


Defined as 1.0 equivalent as the important carbon containing starting material

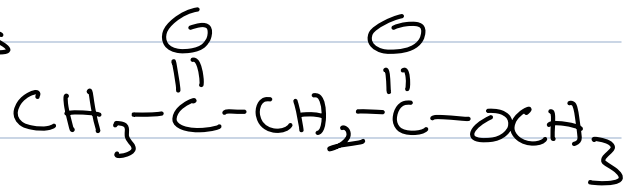
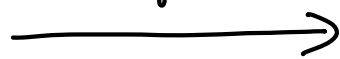
HOCH_3
1.0 equivalent

NaCl
0.5 equivalent

Example of using equivalents



1) NaOCH_3
0.5 equivalents



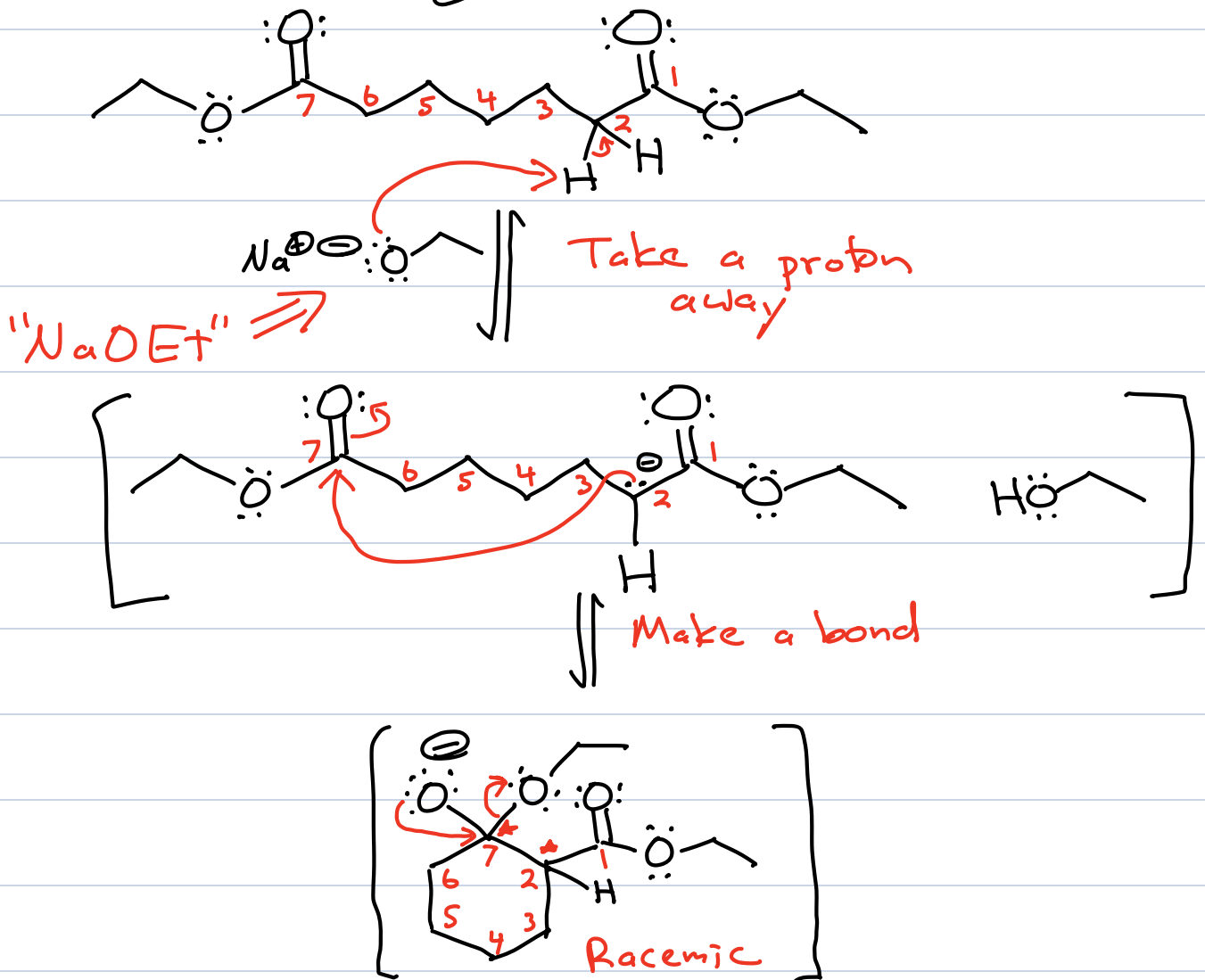
2) $\text{H}_3\text{O}^{\oplus}$

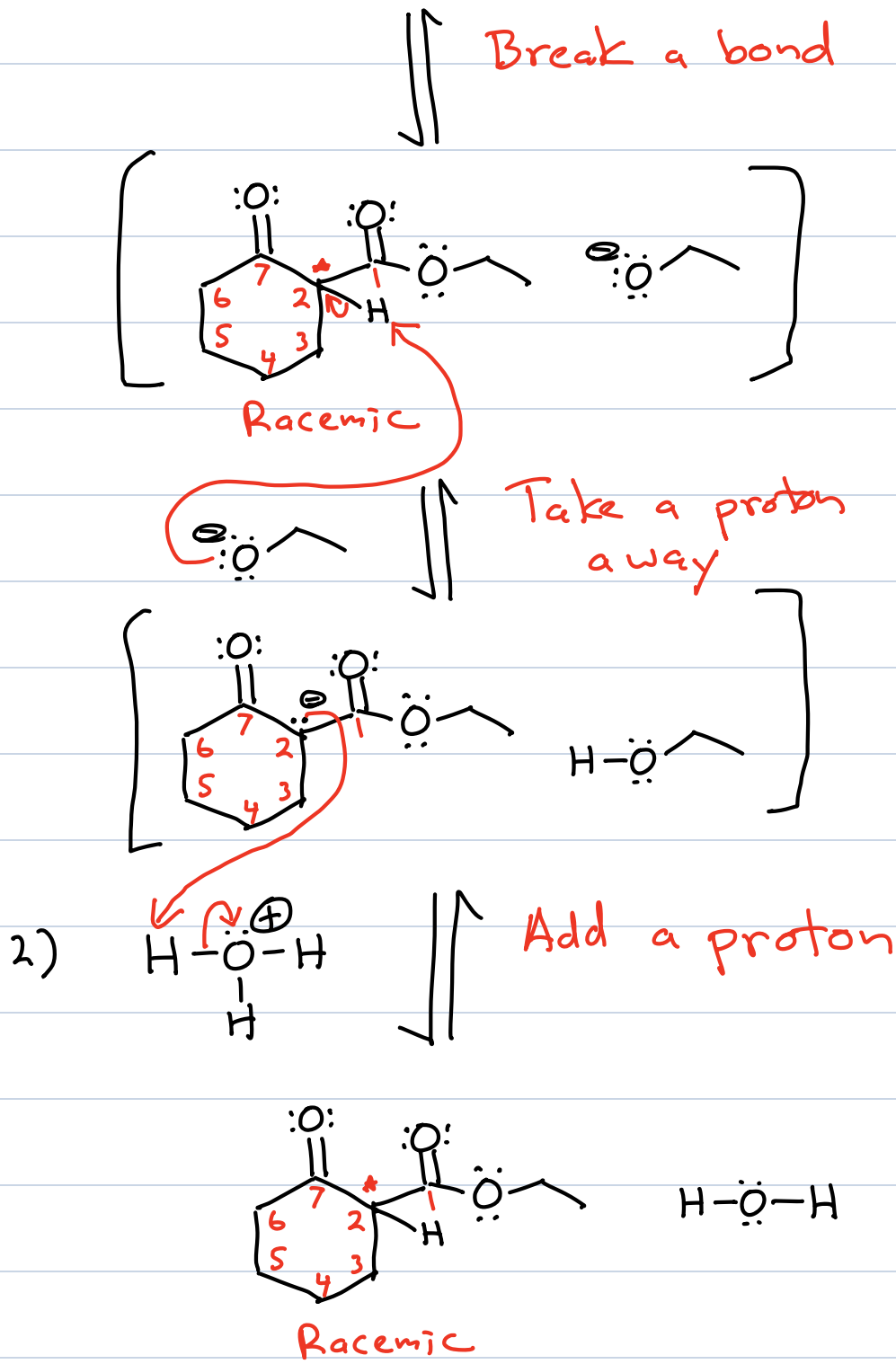
mild

0.5 equivalents
of HCl

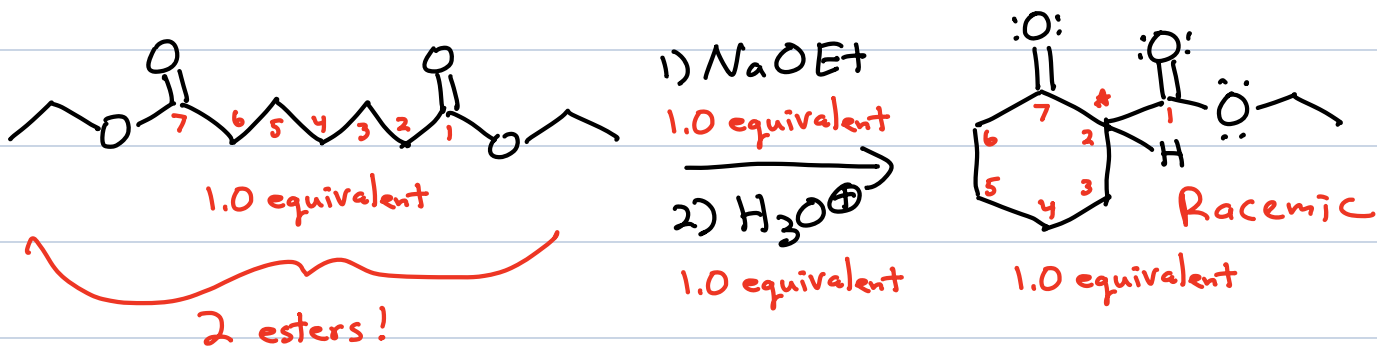


The Dieckmann Condensation →
Using a Claisen to make a 9
ring.

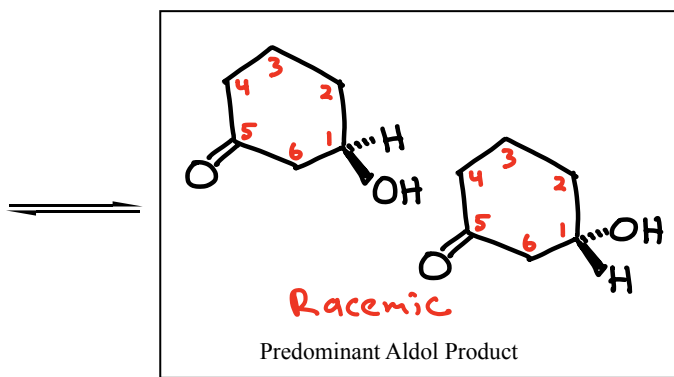
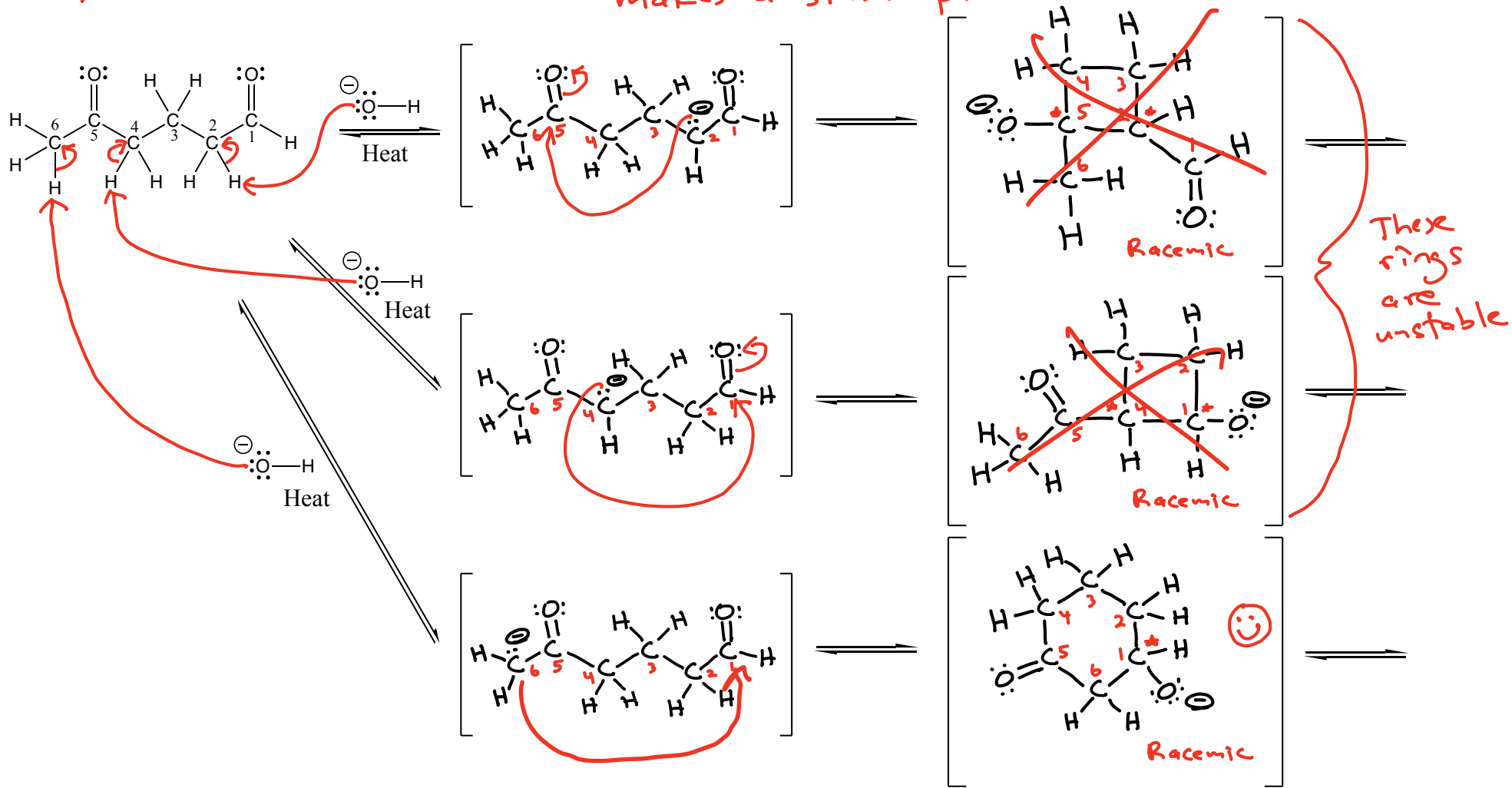




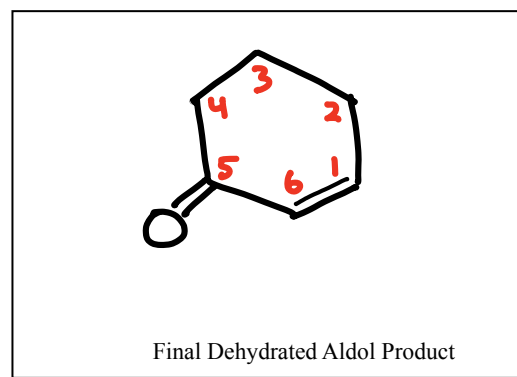
Overall Process



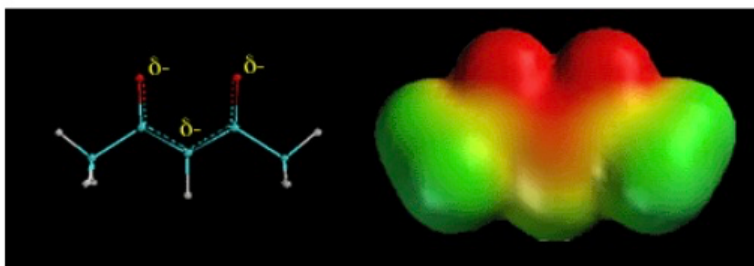
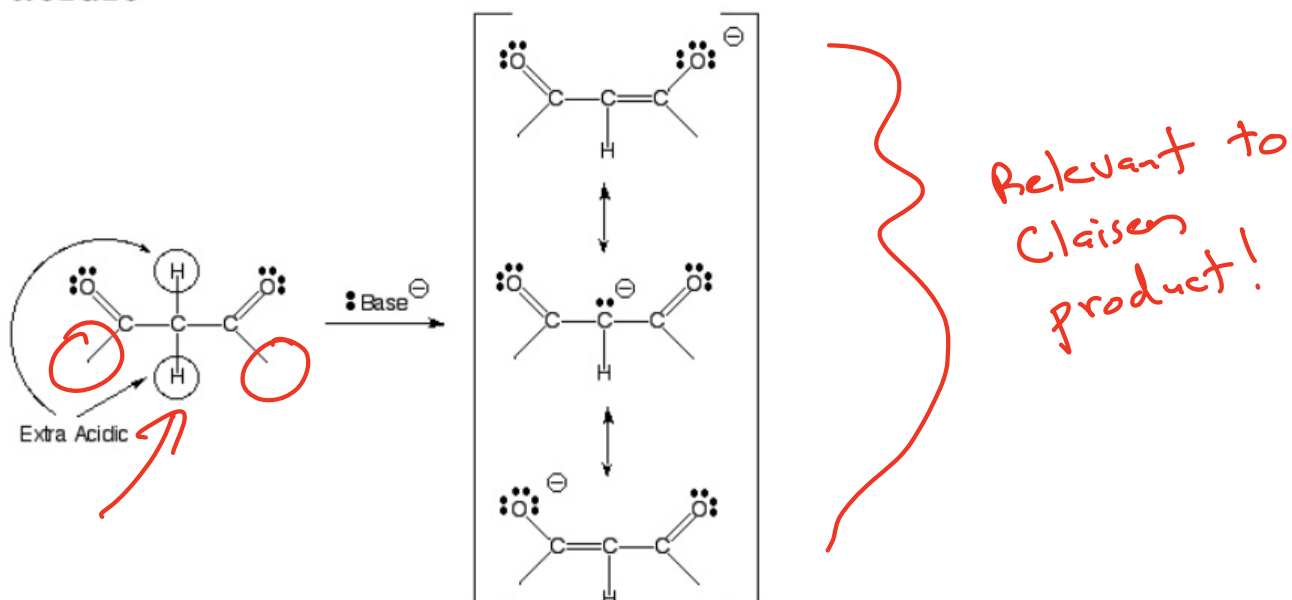
Cyclic Aldol Reaction → 3 different enolates are possible, but only one makes a stable product



Dehydration



Beta-dicarbonyls have alpha-hydrogens that are extra acidic



The C-H hydrogen atoms between two carbonyl groups are even more acidic than normal hydrogens because the resulting anion is double resonance stabilized. The above electrostatic potential surface shows how the negative charge (red color) is spread over all three atoms as predicted by the three resonance contributing structures.

Weaker bases are favored at equilibrium

Compound	Chemical Structure	pK _a
	H-Cl	-7
Carboxylic acids*	$\text{R}-\text{CO}-\text{H}$	3-5
β -Dicarbonyls*	$\text{RC}-\text{CH}_2-\text{CR}'$	10
β -Ketoesters*	$\text{RC}-\text{CH}_2-\text{COR}'$	11
β -Diesters*	$\text{ROC}-\text{CH}_2-\text{COR}'$	13
Water	HOH	15.7
Alcohols	RCH_2OH	15-19
Acid chlorides*	RCH_2-COCl	16
Aldehydes*	RCH_2-CHO	18-20
Ketones*	$\text{RCH}_2-\text{C}(=\text{O})\text{R}'$	18-20
Esters*	$\text{RCH}_2-\text{COOR}'$	23-25
Terminal alkynes	$\text{RC}\equiv\text{C}-\text{H}$	25
LDA	$\text{H}-\text{N}(\text{i-C}_3\text{H}_7)_2$	40
Terminal alkenes	$\text{R}_2\text{C}=\text{C}-\text{H}$	44
Alkanes	$\text{CH}_3\text{CH}_2-\text{H}$	51

Strongest Acid
(Weakest conjugate base)



Weakest Acid
(Strongest conjugate base)

A) Reactions are favored (i.e. have a motive) if they lead to formation of a weaker acid and/or weaker base.

B) Checking pK_a values can predict if a reaction has a motive even if there are other steps besides a proton transfer.

C) Recall that the conjugate base of a stronger acid (lower pK_a) is a weaker base.

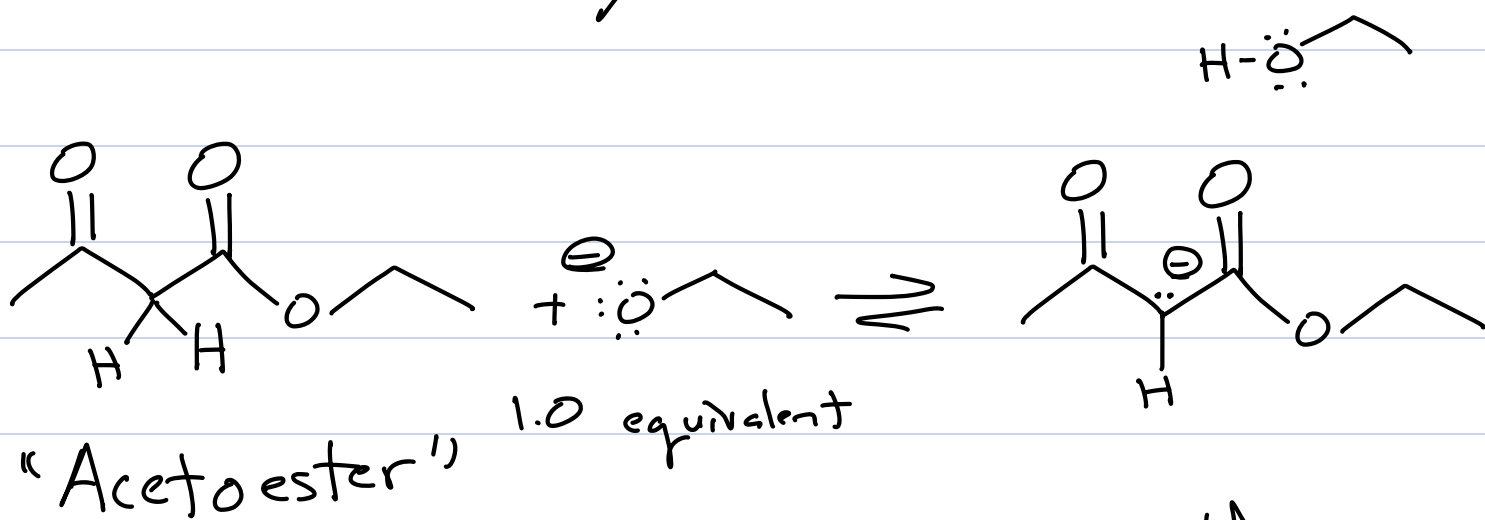
D) Check the pK's of the conjugate acid of the bases on either side of the equation. Lower pK_a value corresponds to stronger acid of the conjugate acid, and thus weaker conjugate base. The base with a stronger conjugate acid (lower pK_a value) will be the weaker base and will be favored at equilibrium.

E) Another way to look at it is that the base that is favored at equilibrium is the one that has the more stabilized anion, i.e. the one with the charge spread around more (electronegative) atoms.

F) Above is a pK_a table that we will refer to often.

*These have resonance stabilized anions

Acetoester Synthesis



$pK_a = 11$

