Recap LDA!

· Kinetic Vs. Thermodynamic Endates

· LDA Allows some reactions that aren't possible w/ Por

(ketone alkylation, crossed aldol, Claisen)

· Reformatsky Rxn:

ROLBY Zn ROLBY

Happy Birthday Colin!

α , β -Unsaturated Carbonyl Compounds: Not Your Typical Alkenes!

Consider the following compound. What functional groups are present? How can we understand the bonding and reactivity of these types of compounds?

How can we explain the mixture of products observed in the following reaction?

α , β -Unsaturated Carbonyl Compounds: Reactivity

In each of the following cases, the "usual" products are observed. What are the products and how are they formed?

Reading: Section 22.8-22.10

α , β -Unsaturated Carbonyl Compounds: Hard and Soft Nucleophiles

Different *types* of nucleophiles tend to favor 1,2- or 1,4-addition. *Hard* nucleophiles are small and have high charge density – they favor 1,2-addition. *Soft* nucleophiles are large and have low charge density – they favor 1,4-addition. But why?!

Hard Nucleophiles	
Alkyl nute lithium	
1 : Ail I.	4 4 h
-OH } but these don't Soft Nucleophiles - 1,4	always give 1,2 -addi.
Soft Nucleophiles ~ 1,4	
R-SH, R-SO	New reagent:
\mathcal{I}^-	New reagent: dialky/ lithium cuprate
R-ZnBr, "soft muleophiles add C'C," Borderline Nucleophiles	R ₂ Cu Li®
"soft muleophiles add ¿ C CI	1
Borderline Nucleophiles	(Culi
NaB Hy	2
-cn mixture	of 1,2, 1,4 addition
RNH2, RRNH mixture	of 1,2, 1,4 addition
\ \\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	2S5
RMgBr)	

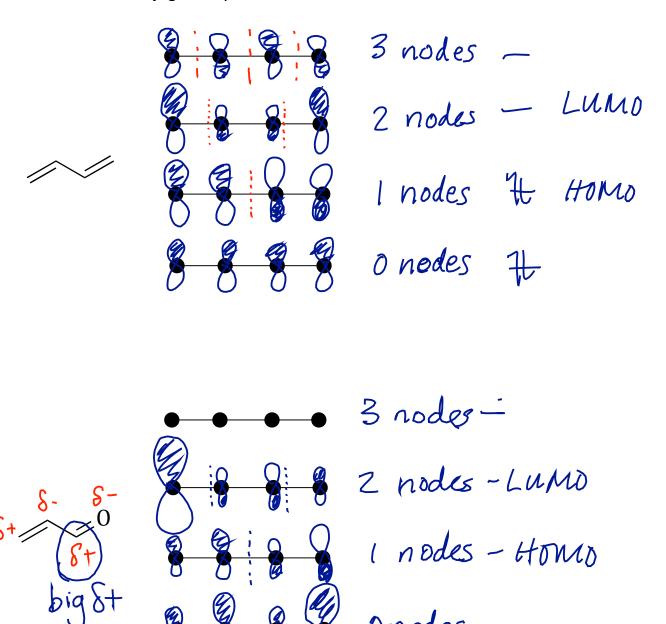
Reading: Section 22.10 KINETIC/THERM CONTROL 10!

for REVERSUBLE NU ONLY

HARD - charge density - charge/polarity
Week 5 SOFT - low charge density - orbital control
July 24, 2014

α , β -Unsaturated Carbonyl Compounds: Molecular Orbitals

To understand why both 1,2 and 1,4 addition are possible, let's examine the molecular orbitals of some conjugated systems.



Reading: Section 22.10

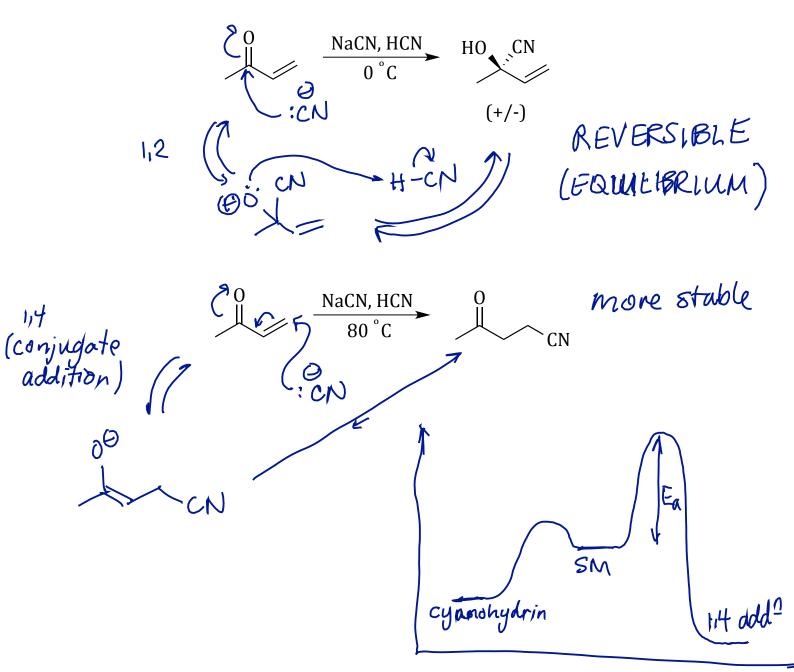
1) Prevent Equilipration! 2) Temperature Week 5

RLi, LiAHY ADD

α , β -Unsaturated Carbonyl Compounds:

Kinetic vs. Thermodynamic Control

Why do the following sets of reaction conditions yield different products?



Kinetic control favors 1,2-addition; thermodynamic control favors 1,4-addition.

Reading: Section 22.10

Why CHz Li 42?

107

IRREVERSUBLE & HARD

α , β -Unsaturated Carbonyl Compounds:

Tandem Addition/Alkylation

"one-pot

How can we carry out the following transformation?

This is an example of a tandem reaction. Why are such reactions useful?

α , β -Unsaturated Carbonyl Compounds: Stereochemistry of Conjugate Addition

Dialkyl cuprates (and most other soft nucleophiles) are very bulky (why?!). Their bulk makes them very sensitive to steric effects in 1,4-addition. How is that fact being used in the following synthesis?

(Made from 2 R/Li+ CUBr -> R2CuLi+ LiBr) FY1 W

109

Arthur Michael

Week 5 July 24, 2014

α , β -Unsaturated Carbonyl Compounds: The Michael Reaction

Propose a complete curved-arrow mechanism for the following reaction. What's going on here?

What kinds of enolates can be used in the Michael reaction?

Michael requires Super soft enclates...

~best dicarbony/enclates

~never w/ lithium enclates (LPA)

How can you recognize the Michael reaction?

1,5 - dicarbonyls

Reading: Section 22.8C

Retrosynth: nore stable enolate Dermabond - octyl cyanoacy late
Week 5

July 24, 2014

α , β -Unsaturated Carbonyl Compounds: Krazy Glue

Tandem Reactions:

The Robinson Annulation ring formation

A remarkable tandem reaction is the *Robinson Annulation*, which is incredibly useful for making 6-membered rings. Provide a complete curved-arrow mechanism:

MICHAE



Retrosynthetic Analysis:

The Robinson Annulation

