

Course 343-lecture 3 Lecturer Gellman  
 Day Friday Date 10-5-12  
 Notes Taken By Matt Aronoff Total # of Pages 4

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Exam #1 locations

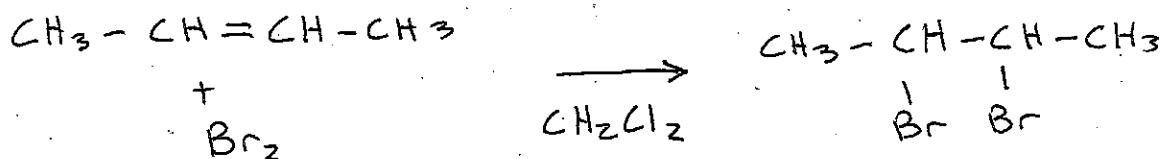
Ackermann - Ji = ~~Ad~~ Noland 132

Jones - Miao = Chem 1315

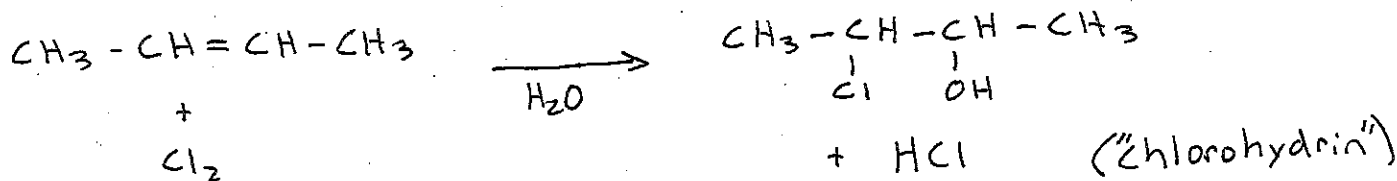
Miller - Zolot = Here (1361)

Recall: Rxns. of alkenes...

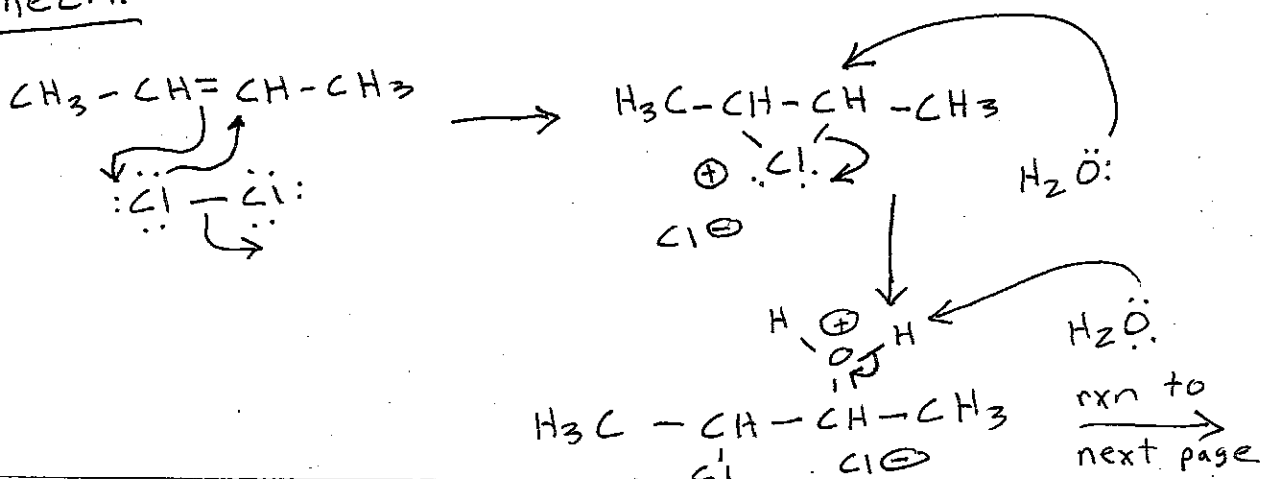
1) w/  $\text{Br}_2$  or  $\text{Cl}_2$  (non-interacting solvent)



2) halohydrin formation

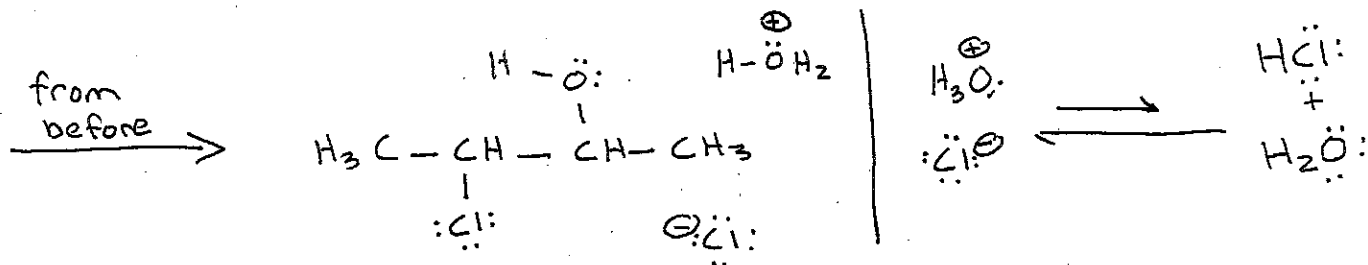


Mech:

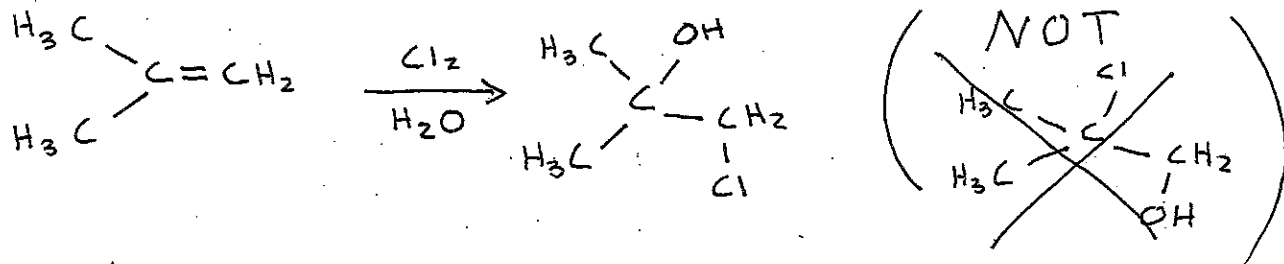


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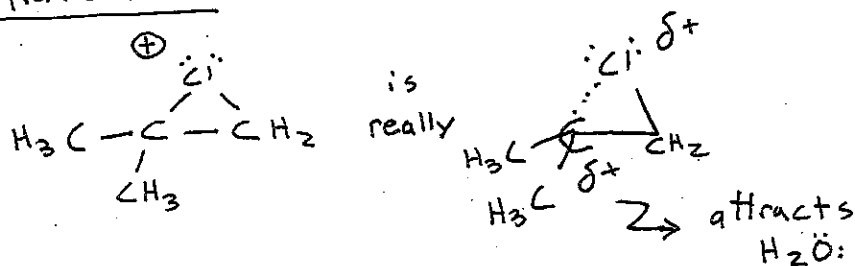
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Unsymmetrical Alkene? Regiochemical Preference? **Yes.**

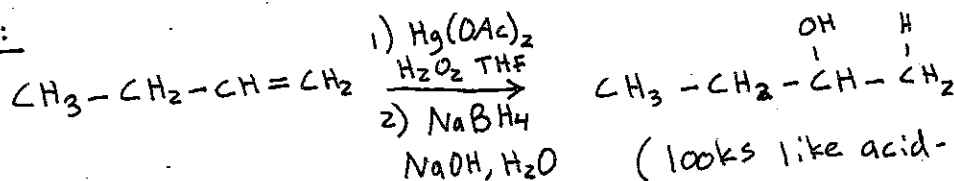


Rationale:



### 3) Oxmercuration-Reduction (2-step process)

Ex:



(looks like acid-catalyzed hydration, but...)

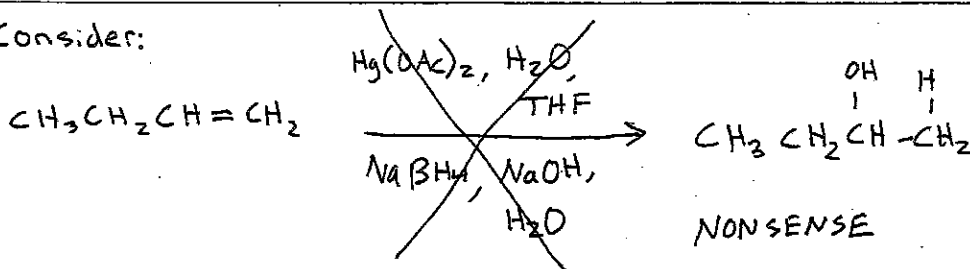
P. 186-  
 use of compact graphical notation to indicate organic transformations

Note:

"1)" vs. "2)" tells us that there are distinct steps

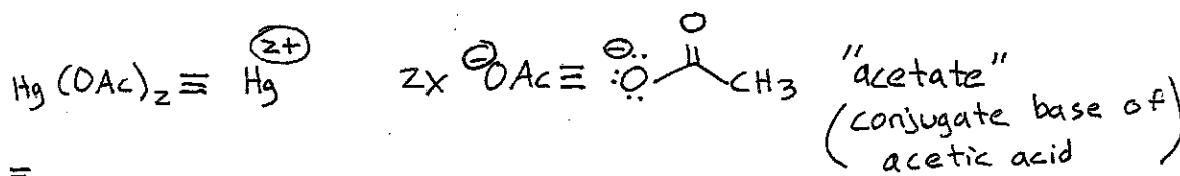
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Consider:

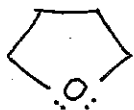


Closer look...

Step #1

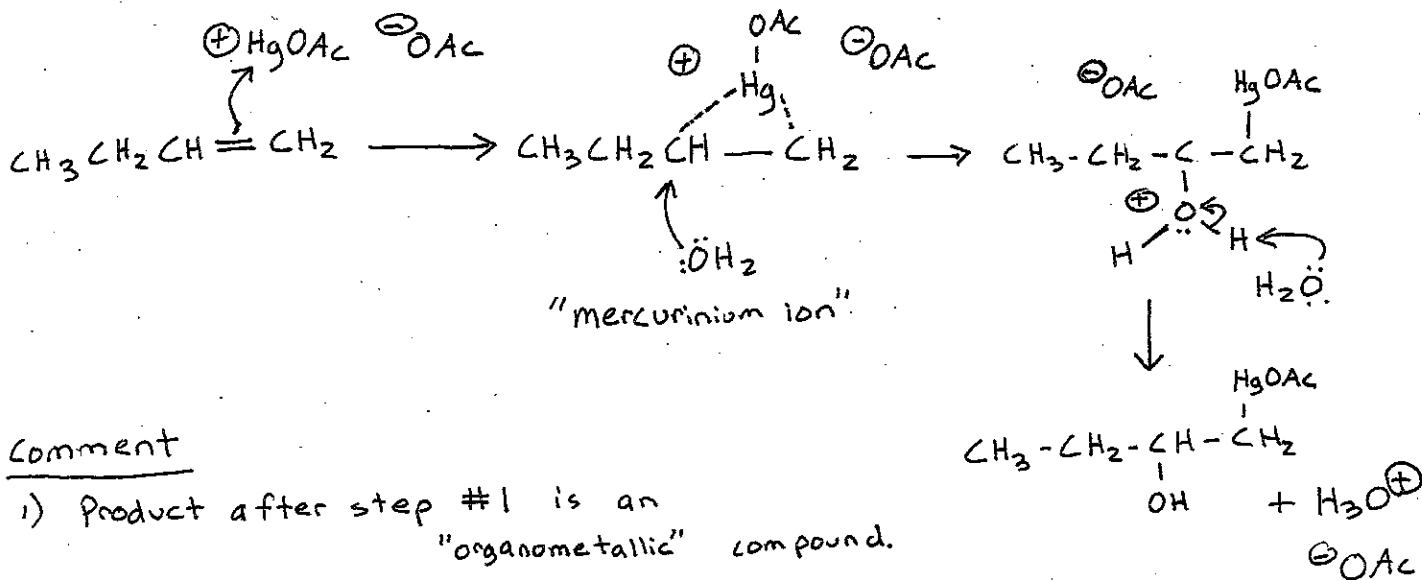


THF  $\equiv$



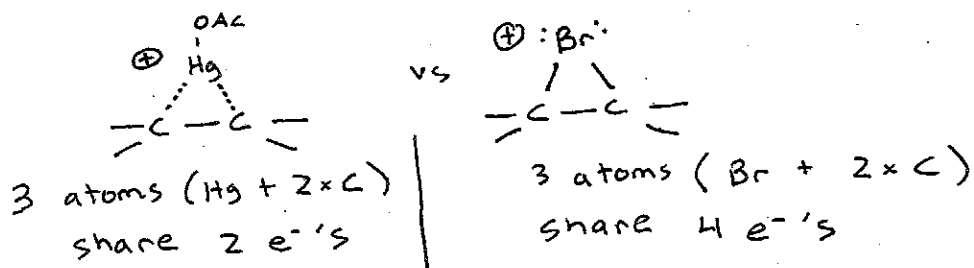
"Tetrahydrofuran"

Mech:



Comment

- Product after step #1 is an "organometallic" compound.
- Nature of the mercurinium relative to bromonium...



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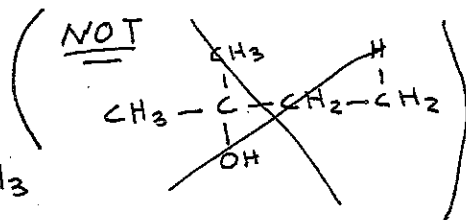
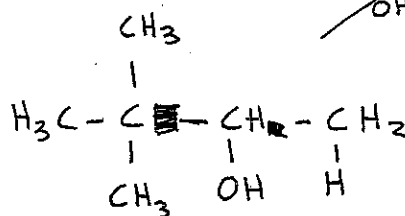
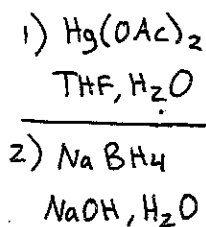
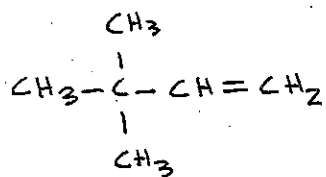
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Shared feature of the intermediates:

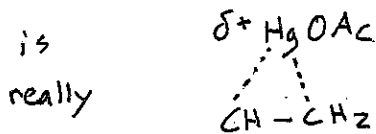
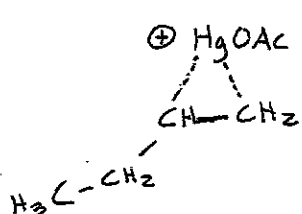
no "free" carbocation — explains why we never see carbocation rearrangements in these rxns.

Ex:



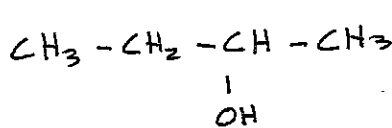
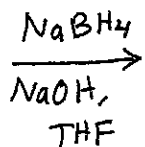
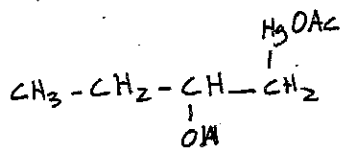
This process gives only the "Markovnikov" product — origin of regiochemical preference?

Rationale:

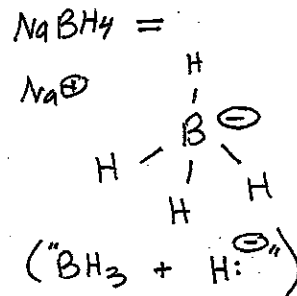


$\text{H}_2\text{O}$  attracted here...

2) Reduction



(+  $\text{Hg}^0$  (↓))



Overall: Accomplish same transformation as alkene hydration but w/o possible rearrangement.

Mechanism not fully understood!