

(36) **Today**

Section 7.7 - 7.11 Electrophilic Addition Reactions

Next Class (37)

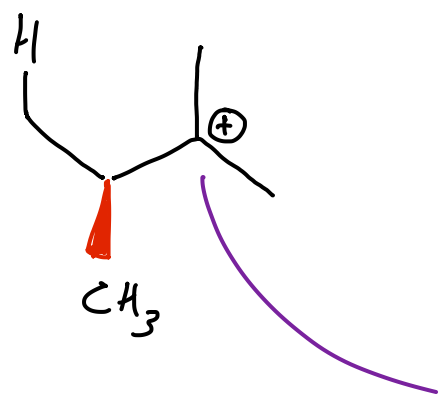
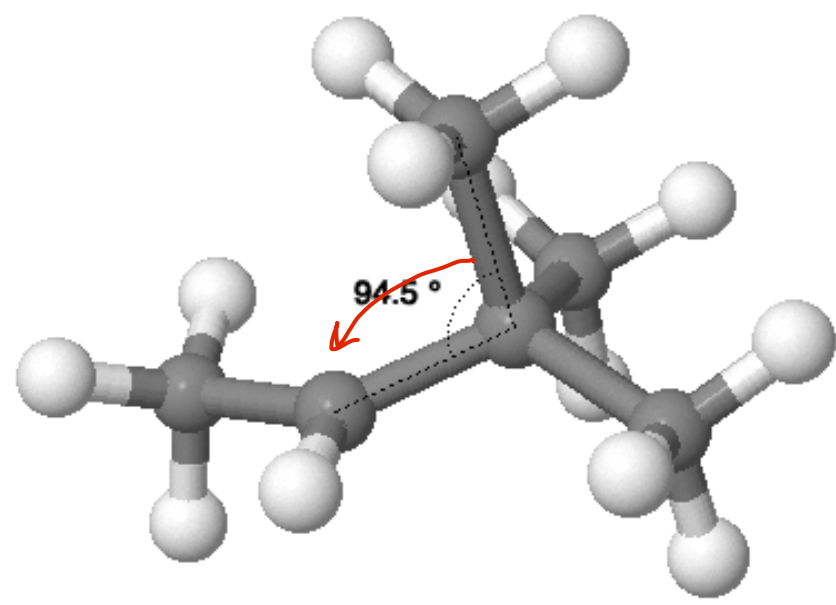
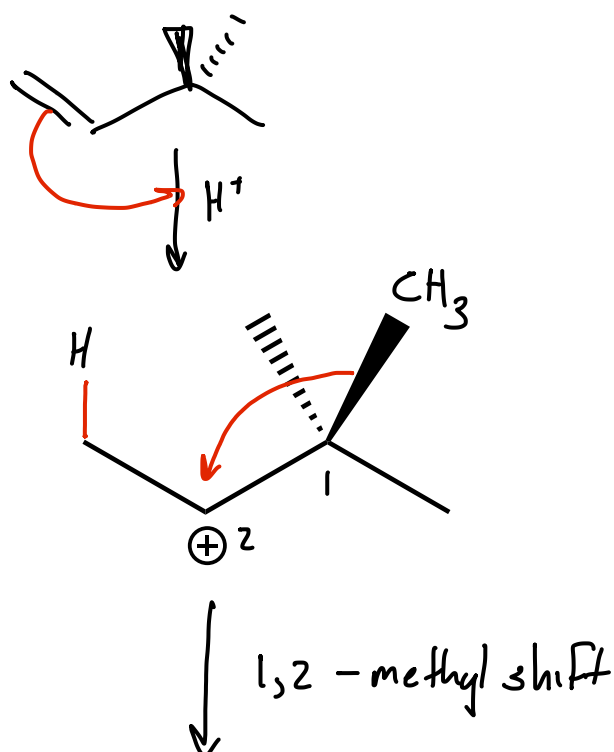
Section 8.2 and 8.3 Halogenation and Halohydrins


the rxns we did in lab

Please Hand in Test Corrections for Test 3 on Your Way Out

Final Is Scheduled for Monday, December 18 from 12:20 to 2:20

The final will be approximately 1 page each from tests 1, 2, and 3, and 1.5 to 2 pages on alkene nomenclature, stability, carbocations, and electrophilic addition.

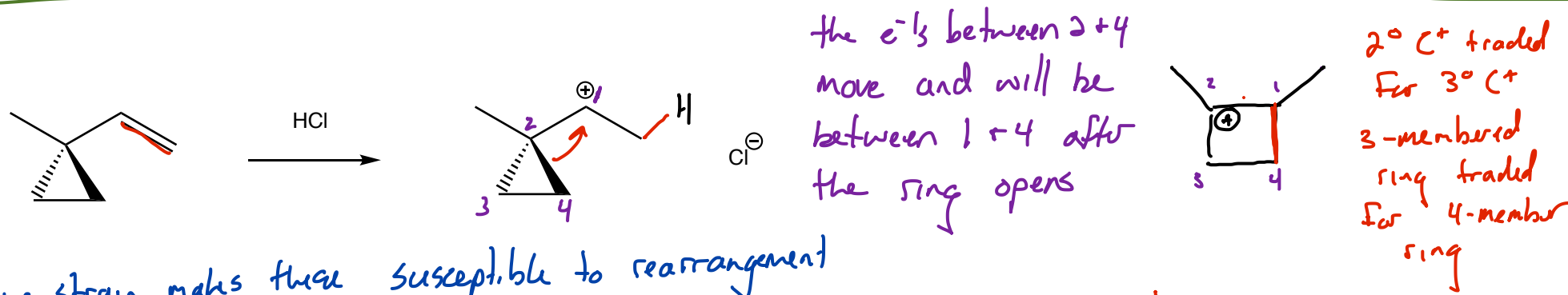
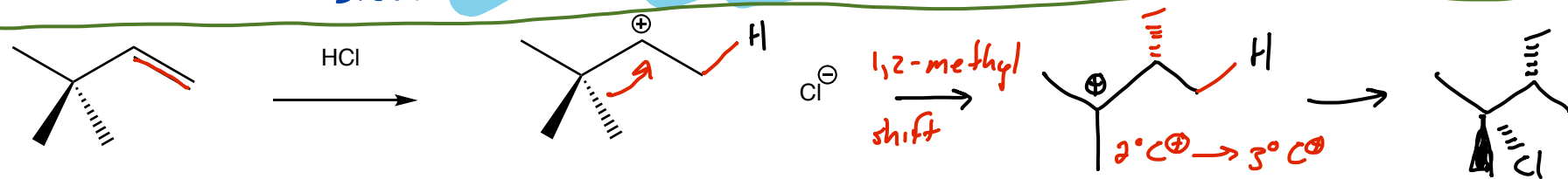
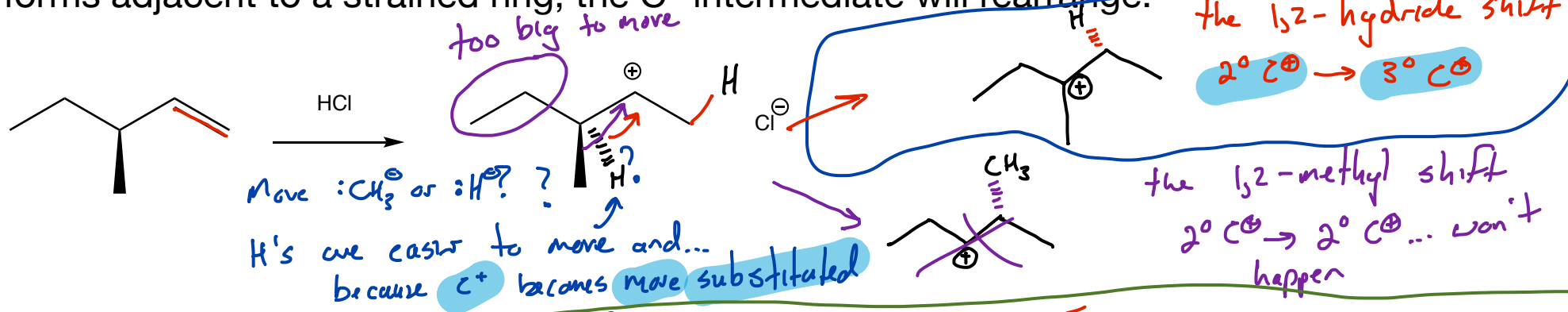


C⁺'s rearrange when they can create a more stable C⁺ on an adjacent C atom
 you may get a small amt. of product from this one
 the rearranged C⁺ is what will react in the 2nd step

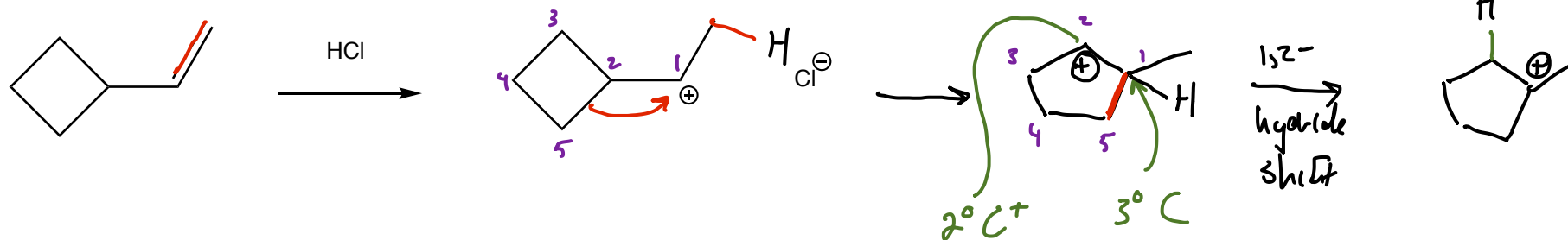
Carbocations rearrange

Section 7.11

If moving an H⁻, or a CH₃⁻, to the position of the C⁺ creates a more stable C⁺ or if a C⁺ forms adjacent to a strained ring, the C⁺ intermediate will rearrange.



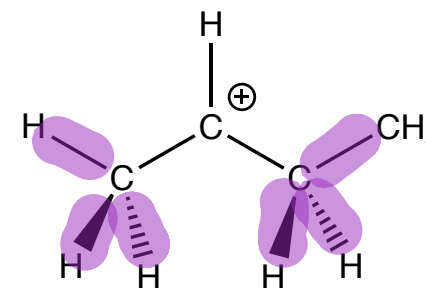
ring strain makes these susceptible to rearrangement



Stability

Getting electron density to a C⁺ helps stabilize the C⁺

e⁻'s in σ-bonds on neighboring carbon atoms stabilize C⁺'s by hyperconjugation



Stability order based on degree of substitution (there are other ways to stabilize C⁺'s that we will see later)



Rearranging C⁺'s

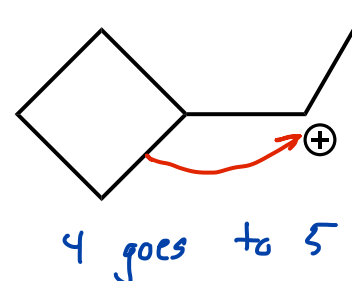
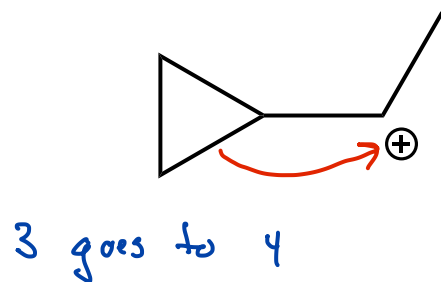
H atoms and methyl groups (CH₃'s) will move from a neighboring C atom if the new C⁺ would be more stable

1° C⁺ will rearrange to a 2° C⁺ or 3° C⁺

2° C⁺ will rearrange to a 3° C⁺

2° don't typically make new 2°... unless

Three- and four-membered rings will pop open if a C⁺ forms adjacent to them.



5, 6, 7... don't do ring opening rxns

~~HF~~

Summary, so far..

Reaction starts at π bond: π bond is lost and σ bonds to electrophile and nucleophile form

Identify the electrophile: so far the H^+ of a strong acid

HCl, HBr, HI

Identify the nucleophile: the conjugate base of the strong acid, the X^- of the HX
an added nucleophile like HOH, CH_3OH, ROH

Create intermediate: open π bond and determine where the $+$ goes (based on stability of $+$)
and attach electrophile to other end.

Are the ends of the double bond identical?

Yes. It doesn't matter; put the $+$ at either end and move on.

No. Is there a reason to prefer making one end of the bond $+$?

Yes. Major and minor products will form. Place the $+$ at the end where it will be more stable and move on.

No. ~1:1 mixture of products will result. Two intermediates form, each one with a $+$ at one end.

Check for carbocation rearrangement: would plus be more stable on neighboring C?

yes, rearrange

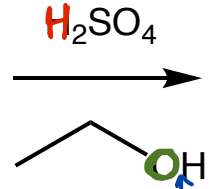
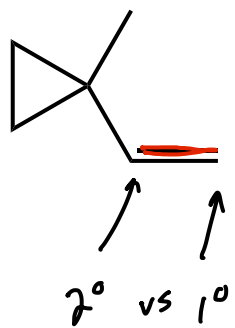
no

leave $+$ where it is

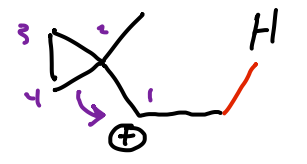
Make a bond from the nucleophile to the C with the $+$ charge.

E Add Reactions

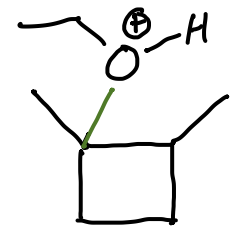
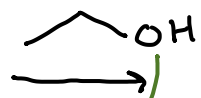
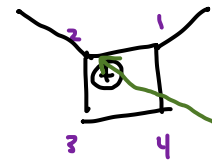
H⁺ of strong acid is the electrophile



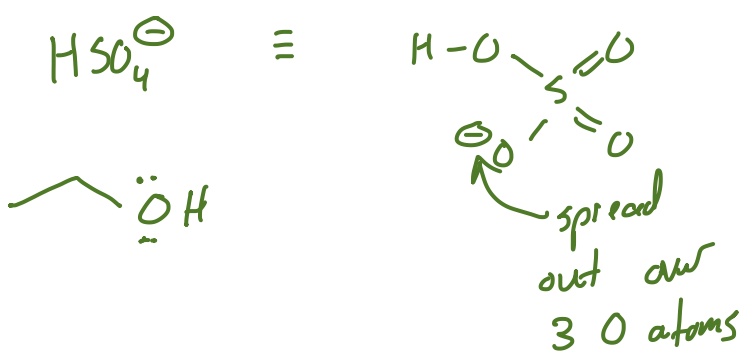
yes polar bond, but not enough like H⁺ of H₂SO₄ is.



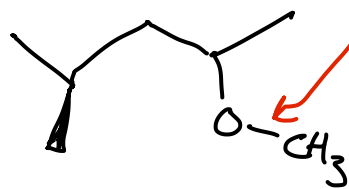
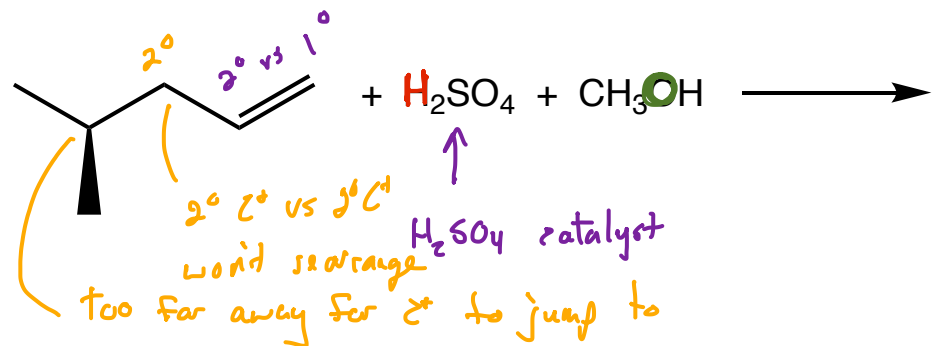
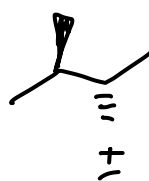
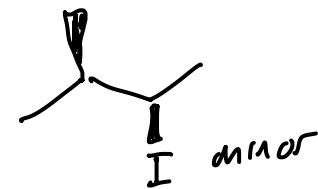
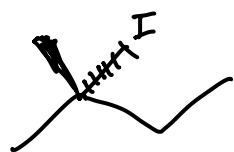
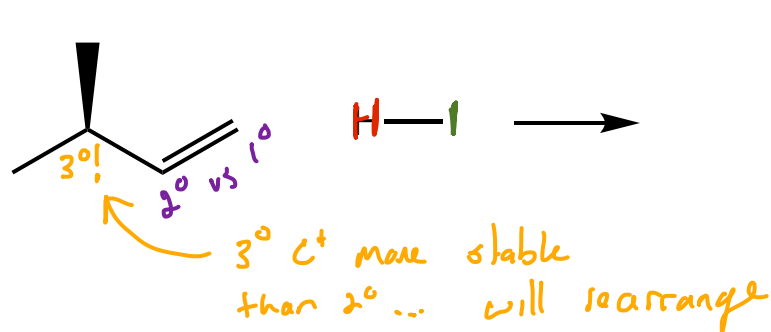
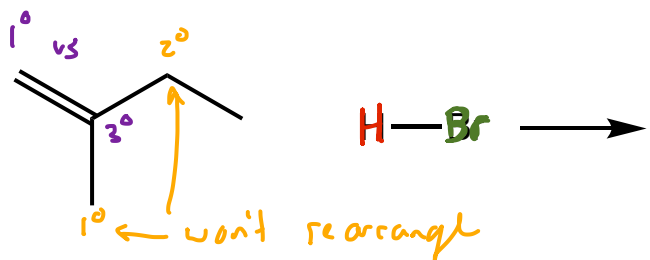
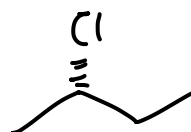
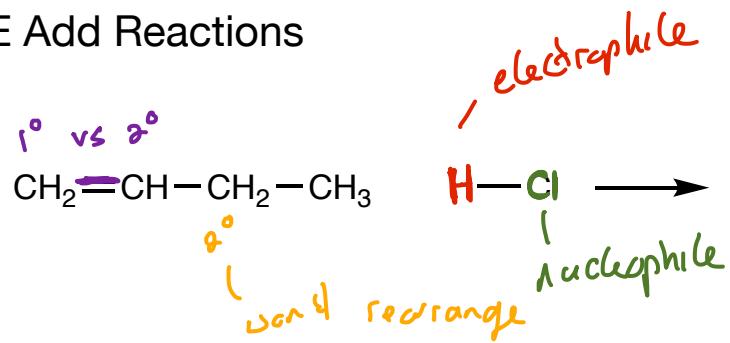
check for rearranging C⁺



H_2SO_4



E Add Reactions



don't break O to C bond
 H_2SO_4

H^+ is lost to regenerate catalyst