

(9) **Today**

Sections 11.7 - 11.11: Practice Elimination Reactions

Competition between S_N1 , E1, S_N2 , and E2

Next Class (10)

Chap 12: Mass Spectrometry and Infrared Spectroscopy

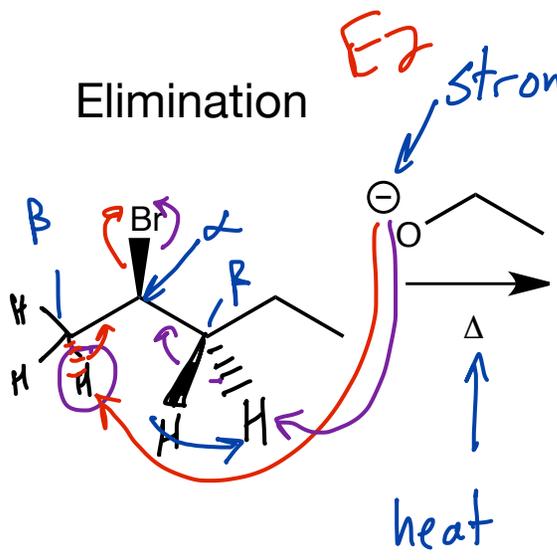
(11) **Second Class from Today**

pseudo-Monday

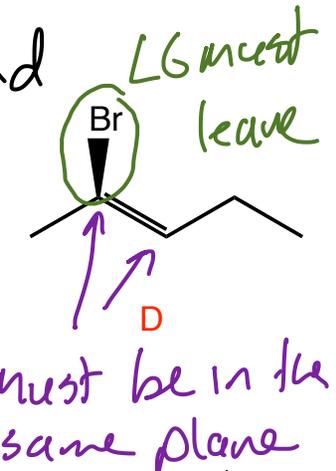
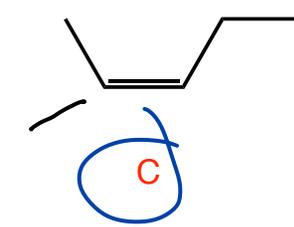
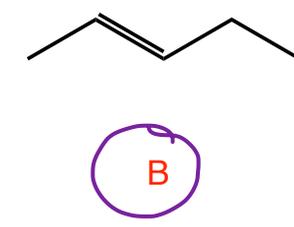
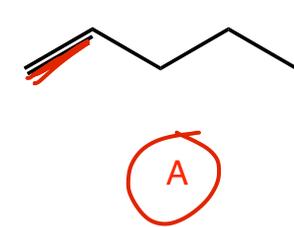
Chap 12: Mass Spectrometry and Infrared Spectroscopy

Third Class from Today (12)

Chap 13: Nuclear Magnetic Resonance Spectroscopy



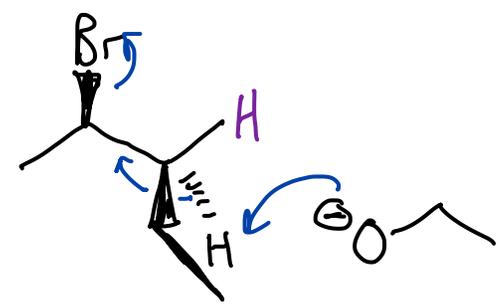
Br is up + in front
 β -H must be down + behind



For the E_2 rxn to occur, the β -C to β -H bond has to be antiperiplanar with the α -C to LG bond

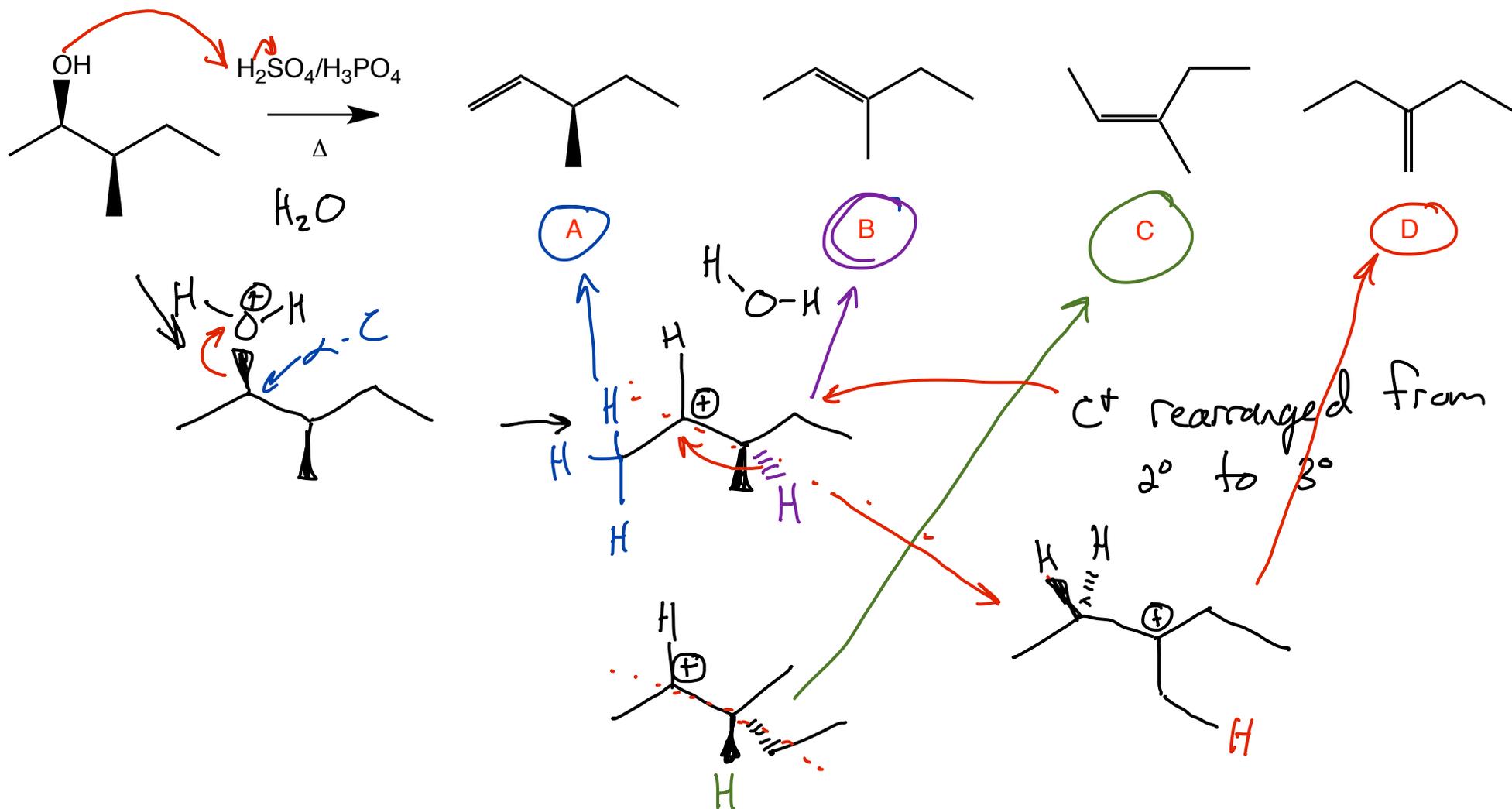
HO solvent
 for rxn
 is typically
 the alcohol
 of the
 alkoxide

C forms when the C-C bond rotates other β -H into position

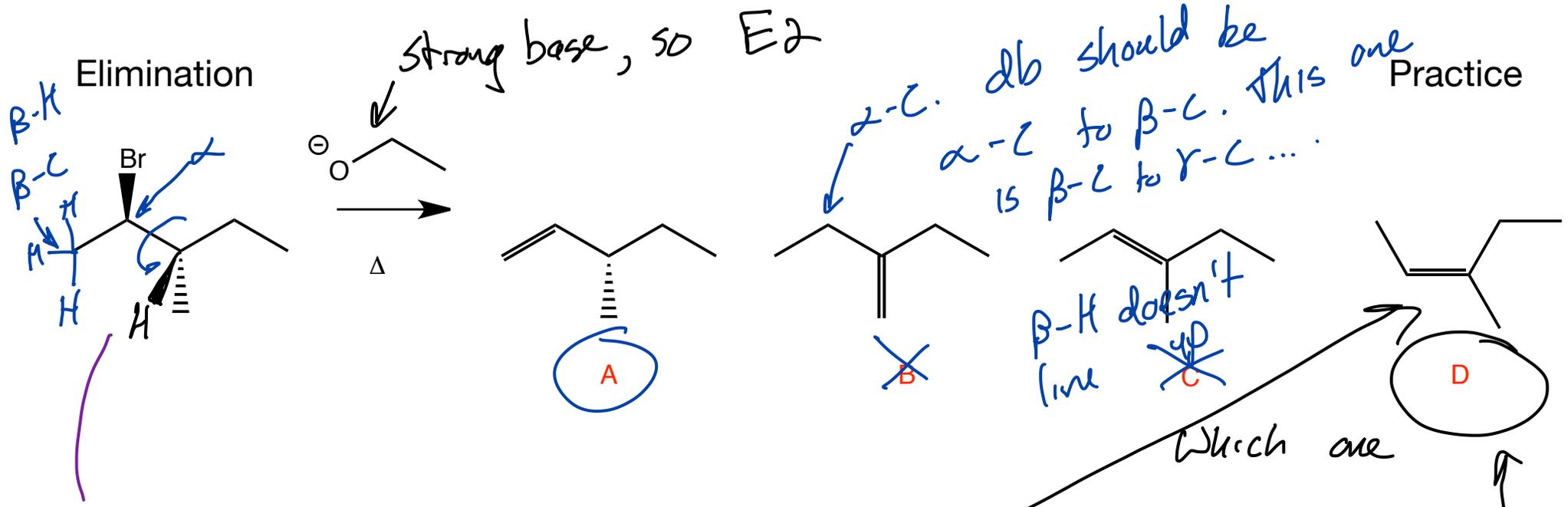


When there is only 1 β -H you have to pay attention to the stereochemical outcome

Elimination E1 Find α - β check for possible C^+ rearrangements Practice



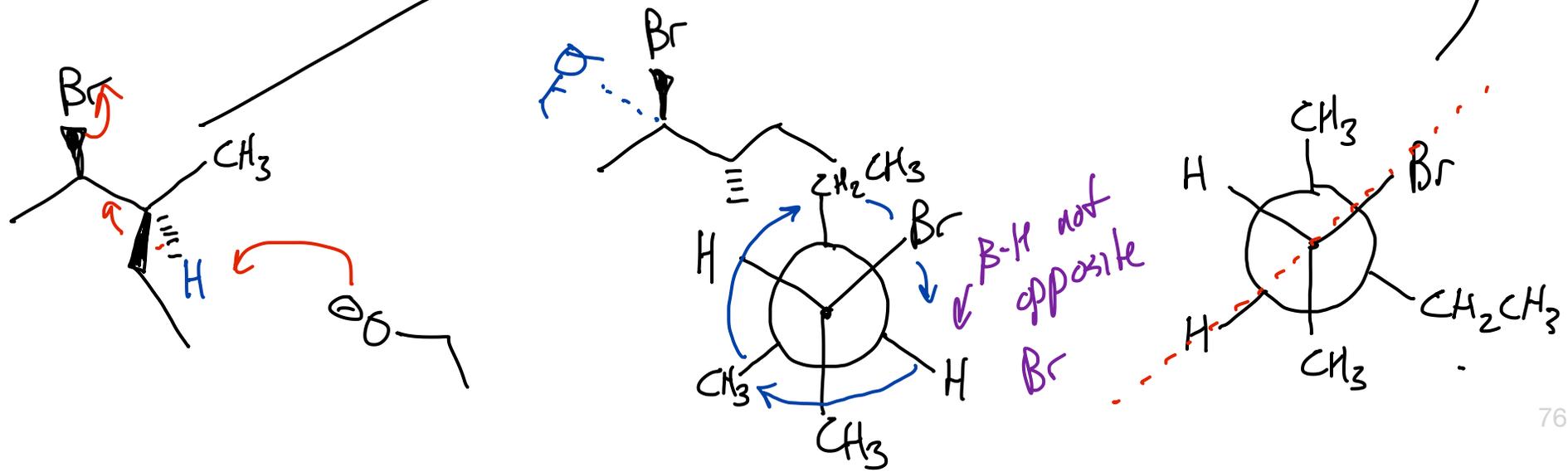
All possible products form. Rearrangements are possible.



this rotamer cannot select to form internal db (C or D)

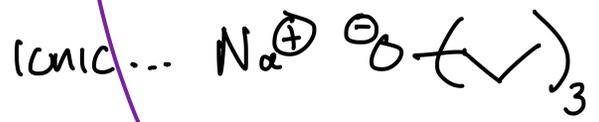
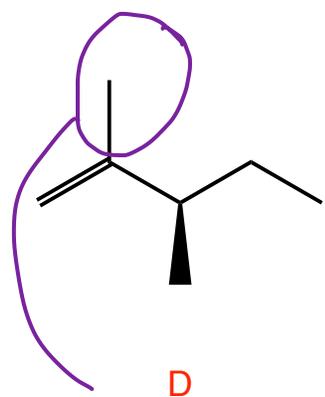
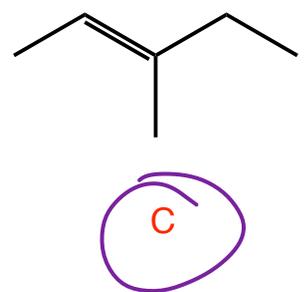
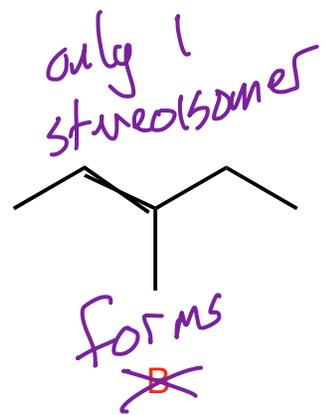
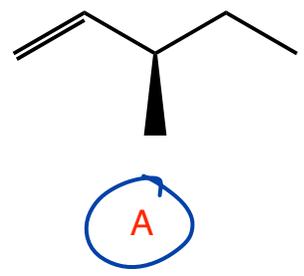
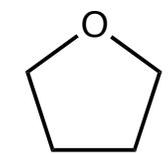
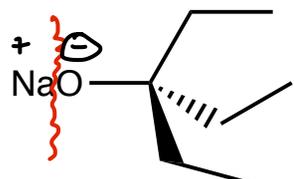
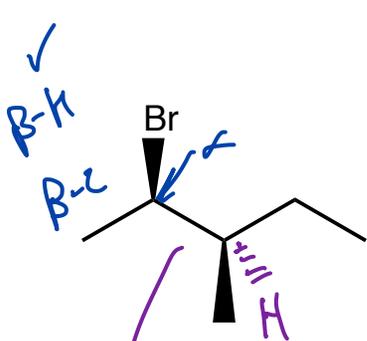
The Br is up + in front and the β -H is down + in front

Need to rotate so β -H is down + behind screen



Elimination Mechanism? E_2 Products? Major product?

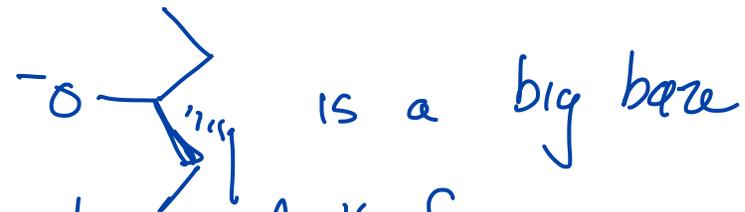
Practice



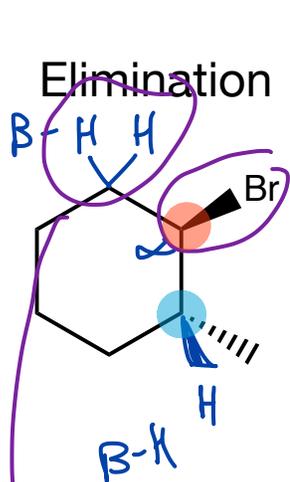
when LG goes the bond goes too

this rotamer has $\alpha\text{-C}$ to LG + $\beta\text{-C}$ to $\beta\text{-H}$
aligned properly

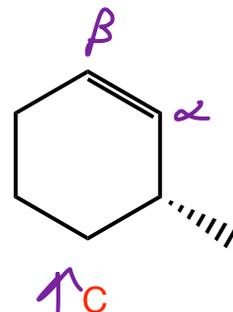
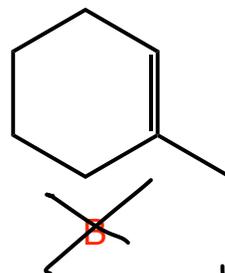
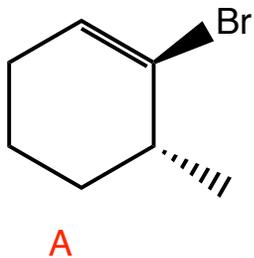
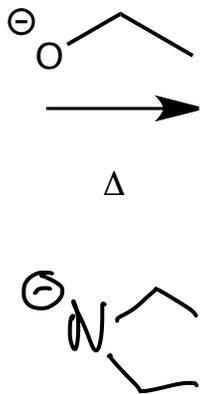
Major product... **A** because



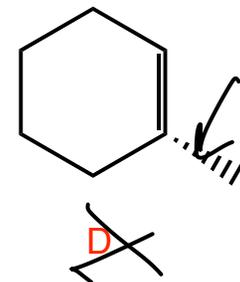
and it will have difficulty abstracting $\beta\text{-H}$ from crowded position



E2

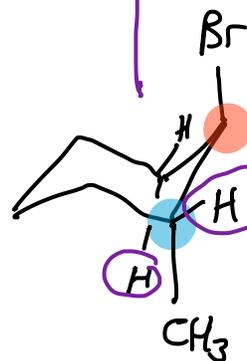
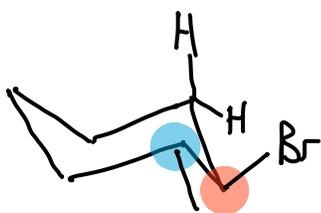
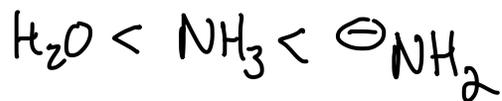
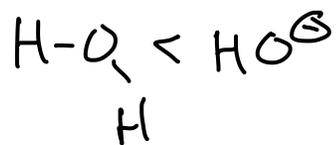


Practice



β -H cannot align to make this prod

Br + one of the β -H can align in a diaxial arrangement



β -H not aligned with axial Br