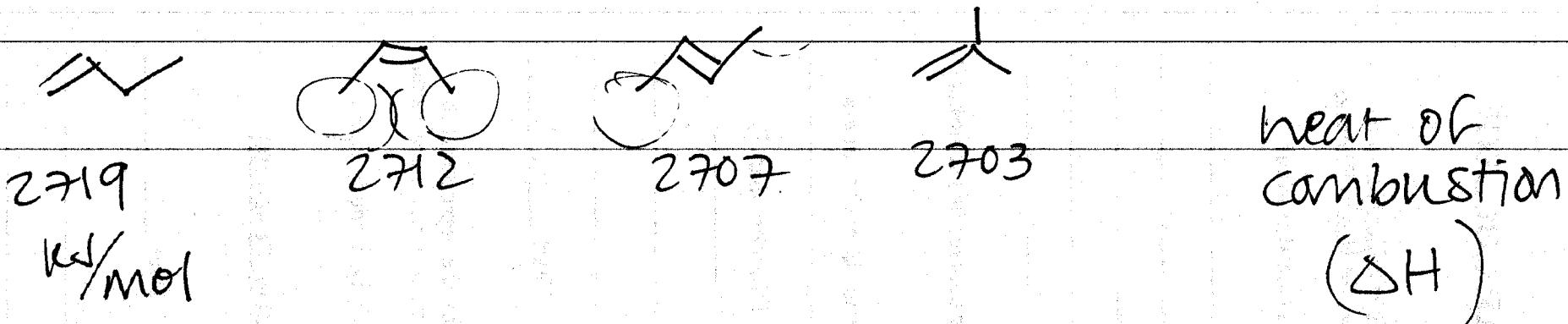


Relative Stabilities of Alkenes



smaller # = more stable

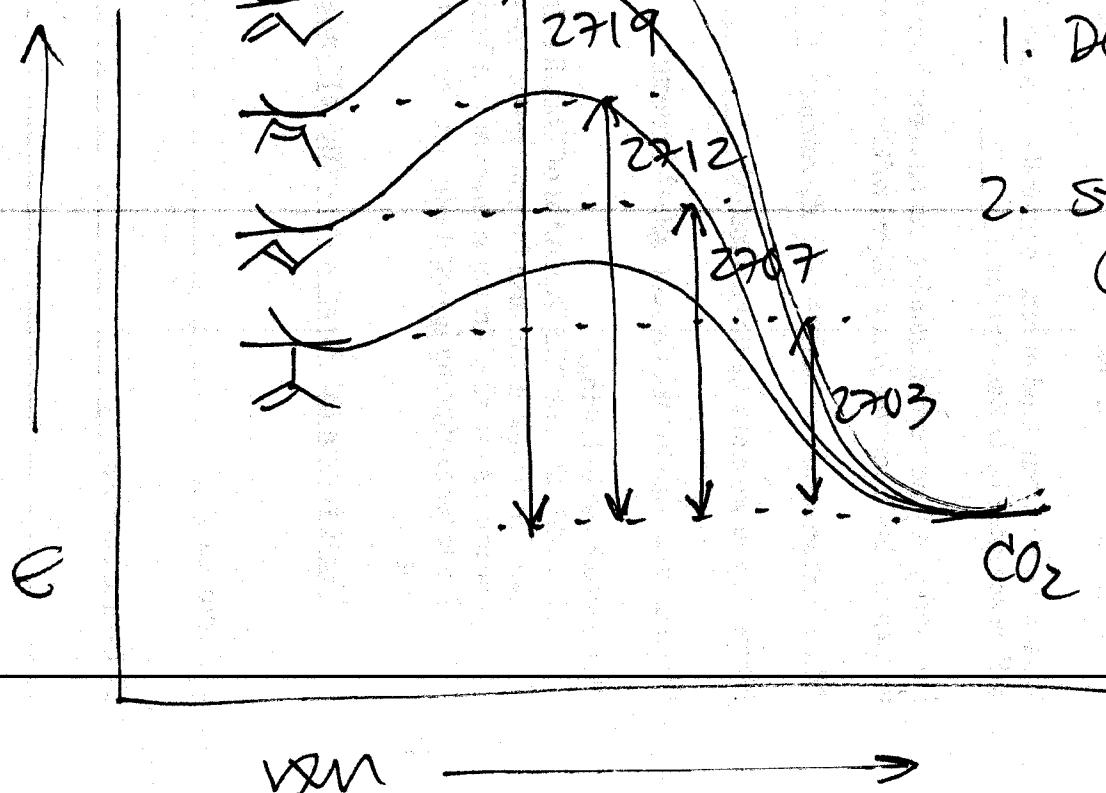
why?

Important factors:

1. Degree of substitution
(more subst = more stable)
2. Sterics

(Z is worse than E)
lower energy

= more
stable



cyclo alkenes - all have \geq double bonds until the ring is ≥ 8 C's.

8-11 C's - can have ϵ alkene in the ring but Z is more stable (ring strain)

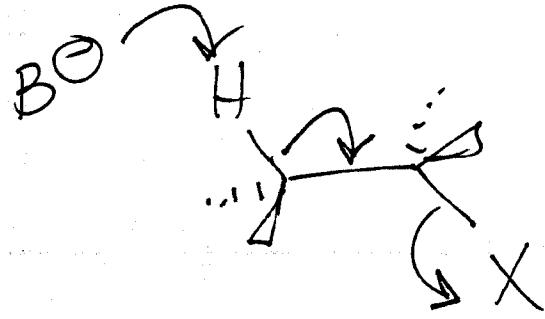
≥ 12 C's - no ring strain associated w/
 Z alkene; ϵ is now more stable

← Don't worry about physical properties.

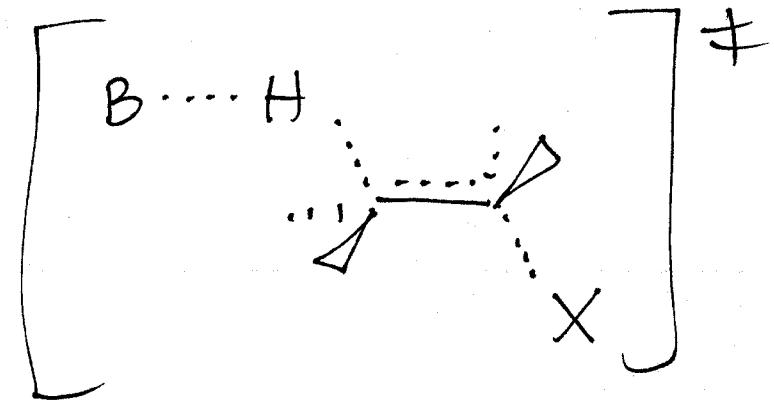
Preparation of Alkenes - elimination Rxn.
* often called β -elim

1. dehydrating halogenation - loss of HX
works best w/ strong base (E2)
worry about: mechanism
regiochem
stereochem. (E vs. Z)

mechanism of an E2: simultaneous ren.



T.S.



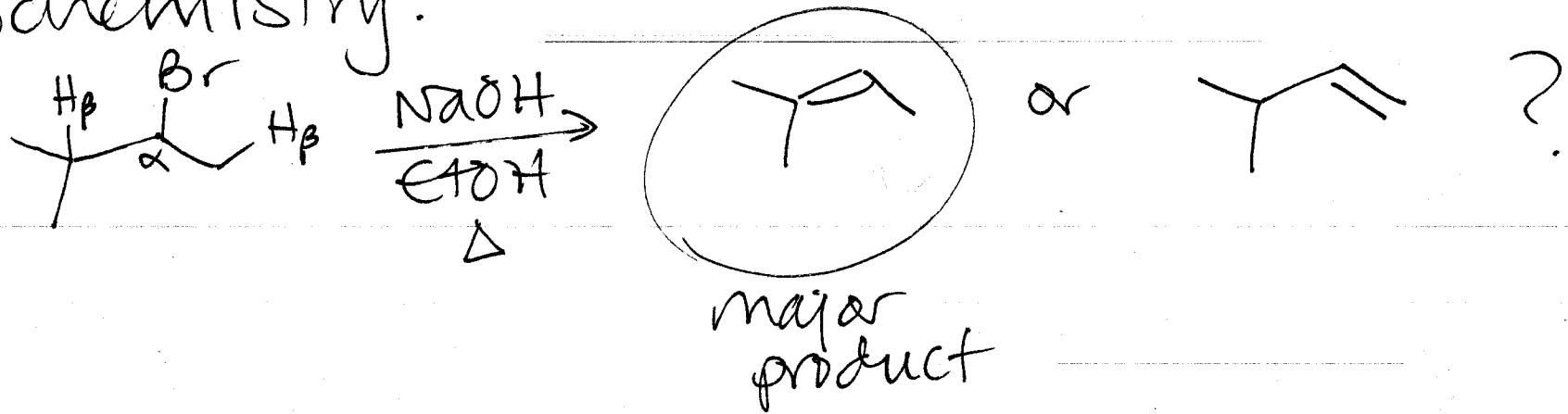
Note: two C's are changing from sp^3 to sp^2 .

How to favor E2:

- $2^\circ/3^\circ$ RX (steric hindrance inhibits S_N2)
- IF using 1° RX use a bulky base ($+OK^+$)

- c. use a strong base (hydroxide or alkoxide)
- d. elevated temp. favors elimination.

Regiochemistry:

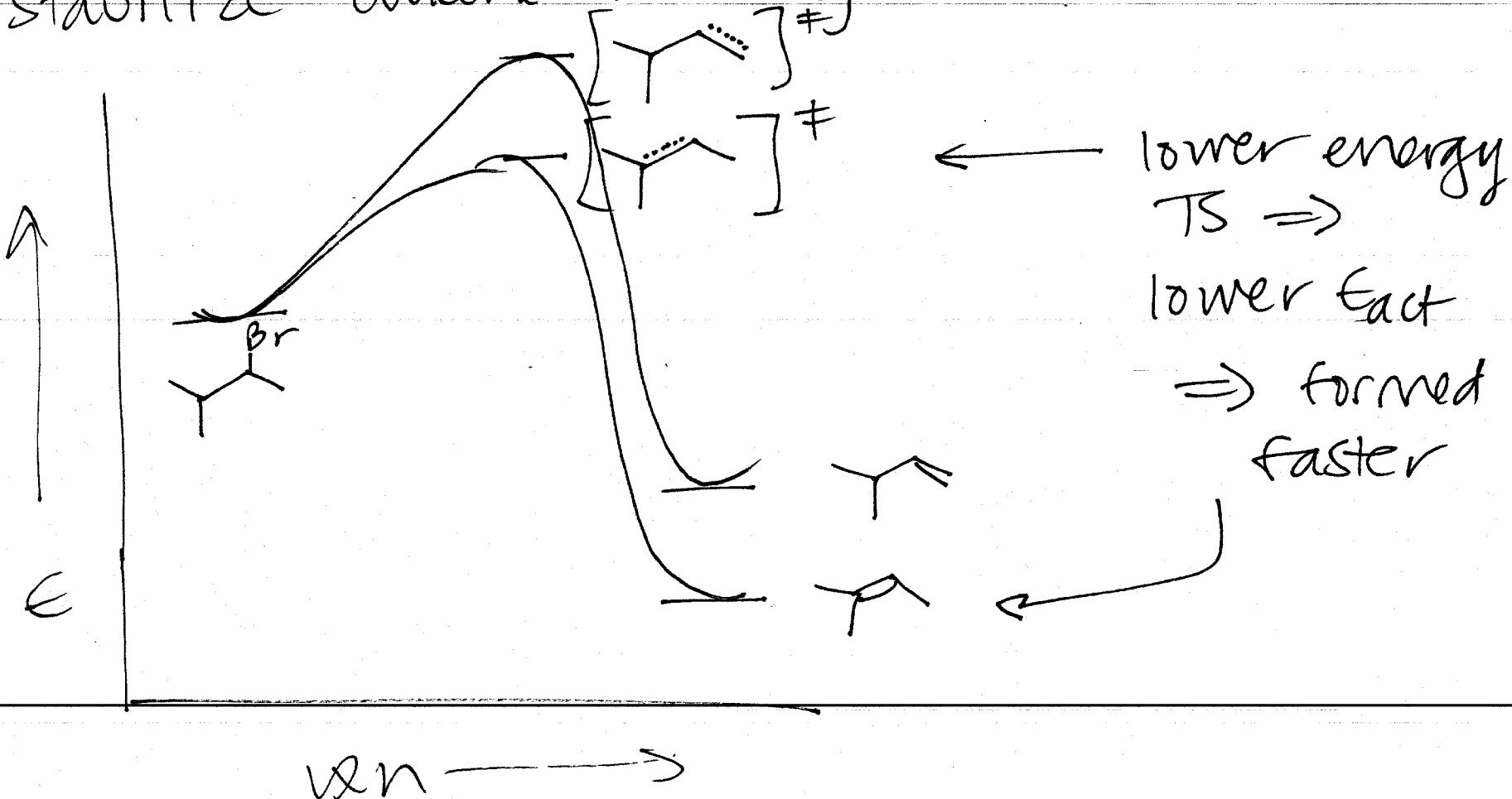


use Zaitsev's Rule: if there is more than one β -H; the one that is lost is the one that will give the more stable product.

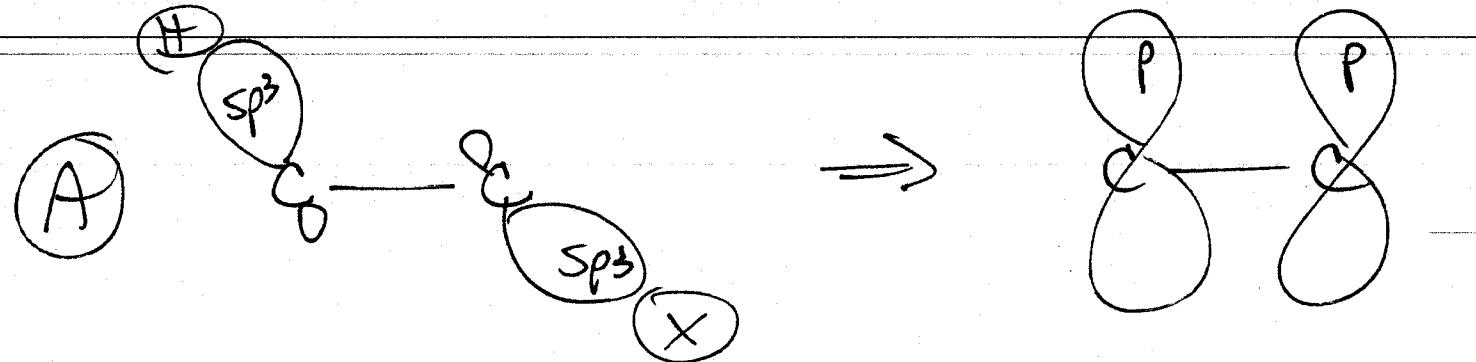
*exception: if using bulky base ($+O^{\circ}K^{\ominus}$) - too sterically hindered to get at 3° H's + has tough time getting at 2° H's - prefers to pull off 1° H's. (Hofmann's Rule - see Ch. 20)

Stereochemistry - ϵ is preferred over γ .

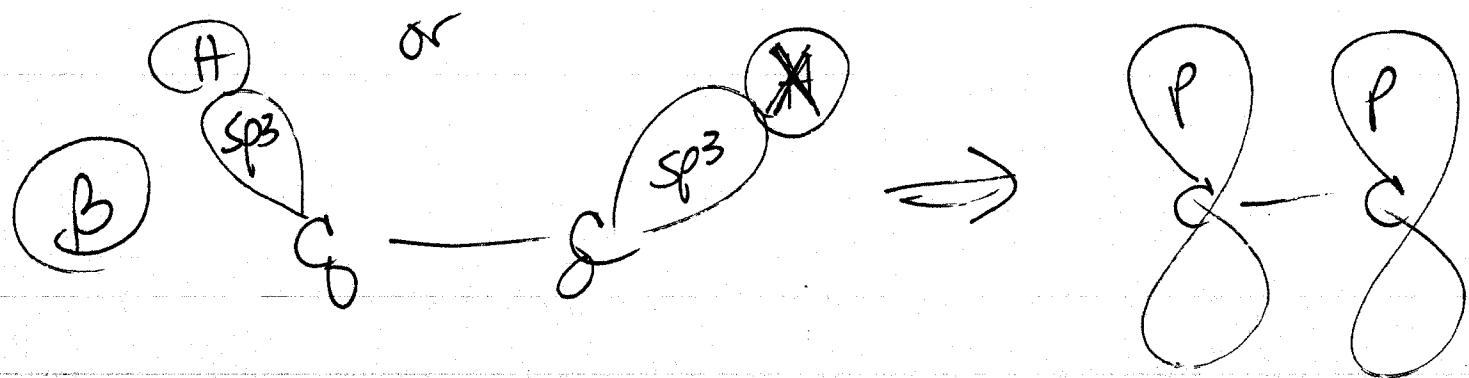
Why does Zaitsev's Rule work? Consider the TS: has partially formed π bond. Alkyl gps. stabilize alkenes; \Rightarrow also stabilize alkene-forming TS.



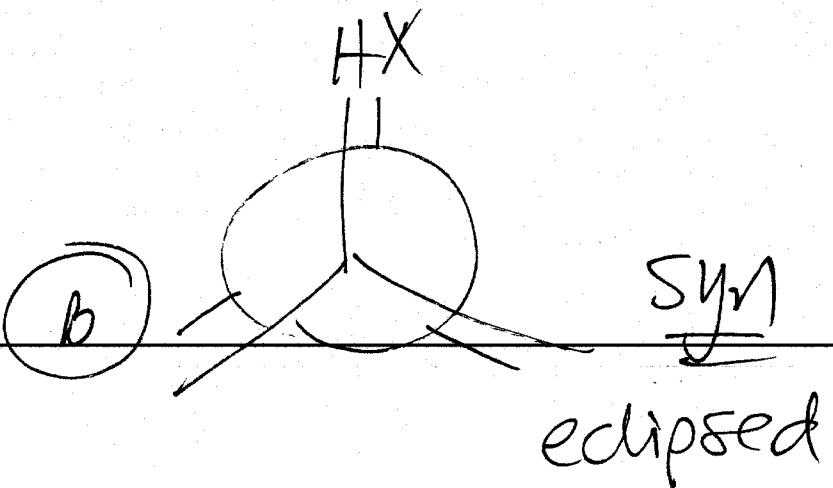
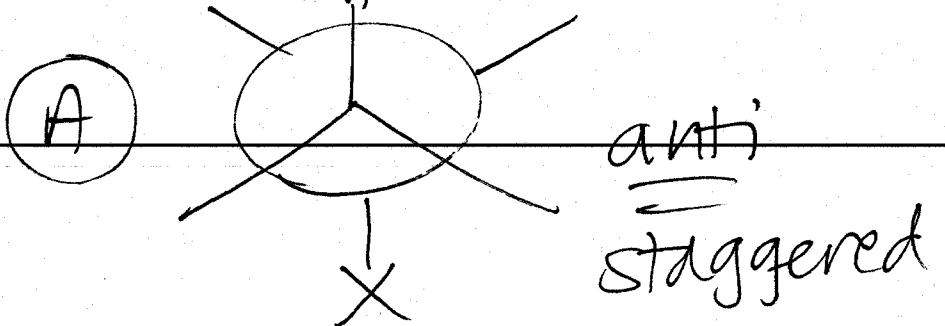
look @ mechanism a little more closely.



p orbitals
need to
be II.



works best when
the two sp^3 's are
already parallel.



consider cis + trans 1-bromo-4-tbutyl cyclohexane

treat w/ HOK/HOH

think about: products + the relative rates.