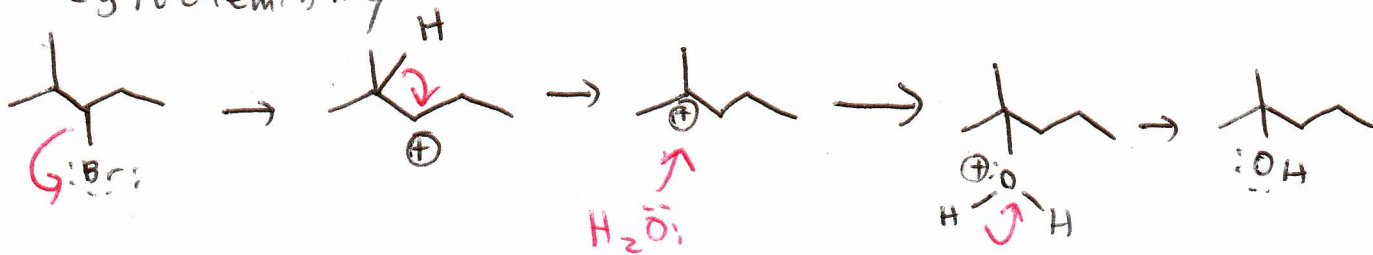


6) Regiochemistry

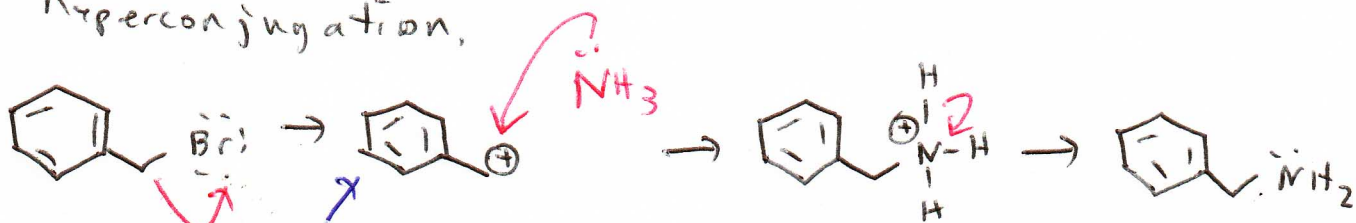


1) substrate

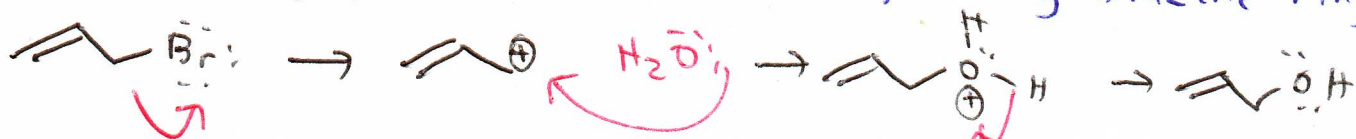
$3^\circ \gg 2^\circ > 1^\circ \sim \text{methyl}$

← reactivity in SN1 reactions

3° substrates are far more reactive in SN1 rxns ~~than~~ since 3° carbocations are far more stabilized by hyperconjugation.



Even though this is a 1° carbocation, it forms easily due to conjugation with the neighboring benzene ring.



Allyl substrates are also reactive due to conjugation.

vinyl C=CX vinyl → attached to the double bond



Vinyl substrates do not undergo SN1 rxns since

vinyl carbocations are unfavorable due to increased s-character of the orbital (sp hybrids better tolerate \ominus charges but poorly tolerate \oplus charges).

Vinyl substrates do not undergo SN2 rxns because the electrons in the π bond repel the nucleophile.

2) nucleophile

For S_N1 rxns, poor nucleophiles - particularly non-basic ones - are preferred, since S_N1 rxns depend on the carbocation having time to form

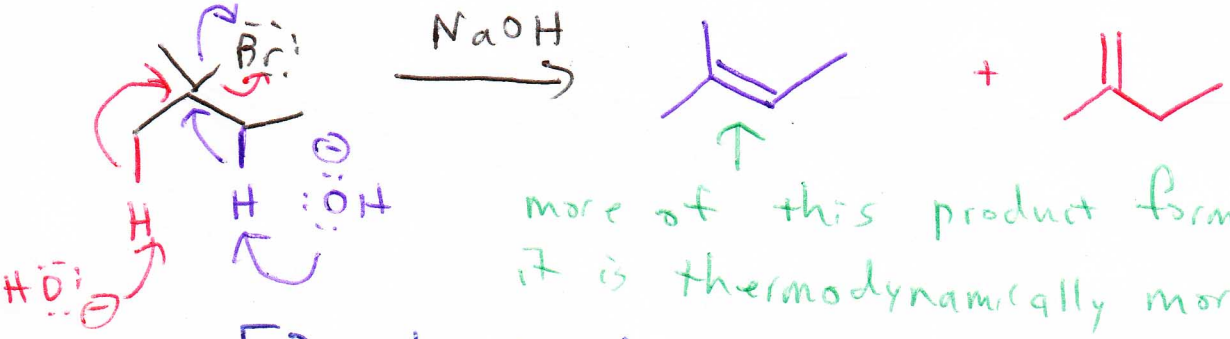
3) leaving groups

Good leaving groups are conjugate bases of strong acids

4) solvent

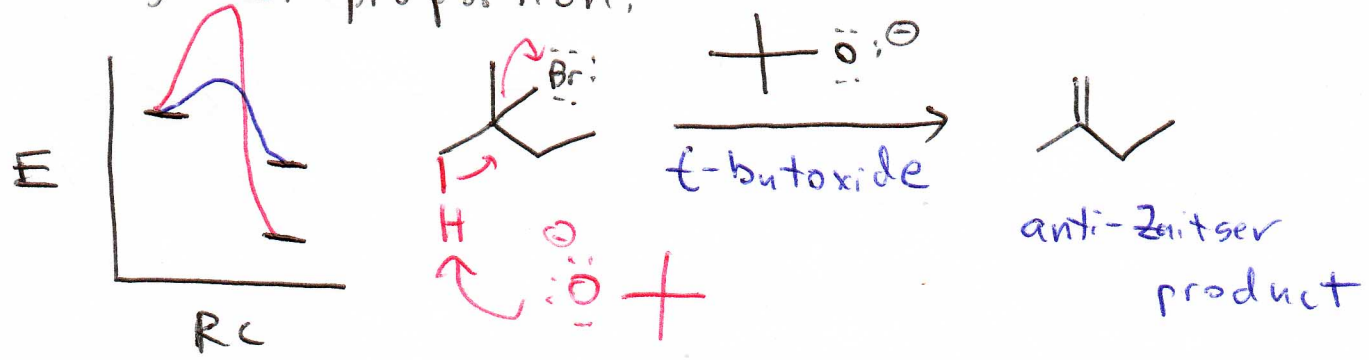
For S_N1 rxns, polar protic solvents are normally used; the nucleophile is often the solvent itself.

Eliminations



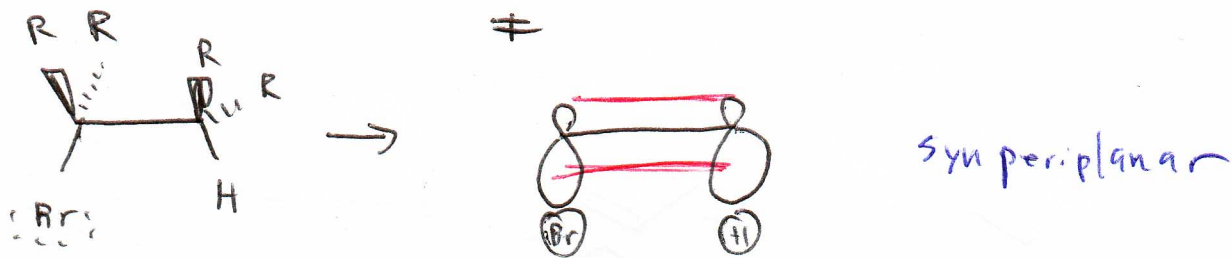
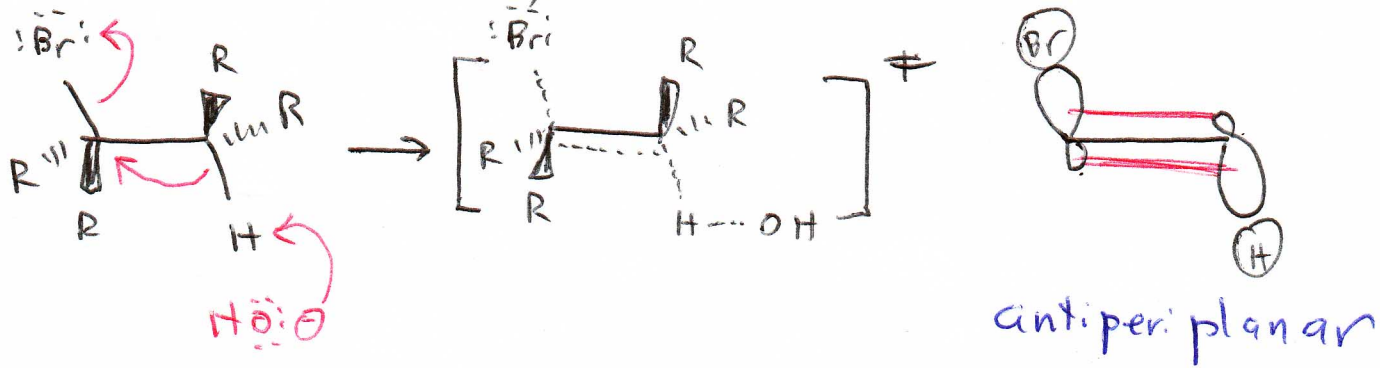
E2 - bimolecular elimination

Zaitsev's rule - In eliminations, the more heavily substituted alkene will normally be formed in greater proportion, *



If a nucleophile is very sterically hindered, it can cause the formation of a less stable alkene due to kinetic effects (more hindered nucleophiles make it more difficult to remove more hindered hydrogens).

- E2
- 1) utility \rightarrow alkyl halides \rightarrow alkenes
 - 2) reagents \rightarrow strong, basic nucleophiles
 - 3) conditions \rightarrow polar aprotic solvents
 - 5) Stereochemistry



Eliminations can only occur if the hydrogen removed is syn-periplanar or anti-periplanar to the leaving group.

