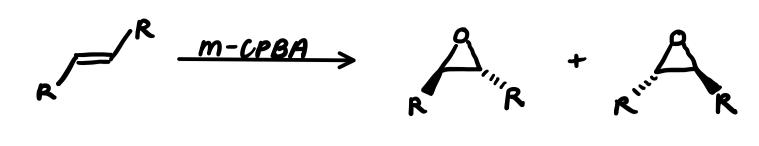
Chapter 11 - Summary of Reactions

Thursday, August 6, 2020



$$R = \frac{i) Br_{3,1}H_{20}}{ii) NANH_{2}} + R^{iii}R + R^{iii}R$$

$$\sim$$
 OH \sim H₂SO₄ \sim

$$\downarrow_{0} \qquad \xrightarrow{HI} \qquad \downarrow_{1}$$

$$\frac{R_3S_{i}-C1}{pyridim} \rightarrow 0 - S_{i}-R$$

- Concerted rxn mah

- alkene geometry preserved (cis-) cis, trans -> trans)
 racemic mixt. if Chiral center formed
- despite going through a diff ryn mech, the stereochemistry gives the same result as m-CPBA
- Williamson ether synthesis RX must be 1° (otherwise EZ)
- ROH must be 1° or 20 (other wise E2)
- ocid cat. ether synthesis
- alkene must be able to form a carbocation

- 1st cans, halogen goes to other carbon that forms the most stable cation
- Stureochem. considerations applicable to SNI, SN2 apply depending on mech.
- HBr also works
- Protecting group for alcohols R= CH3, Others
- removal of protecting group

Epoxide-opening (BASIC/NU)

$$\frac{i) : Nu}{ii H_S 0^+}$$
HO Nu

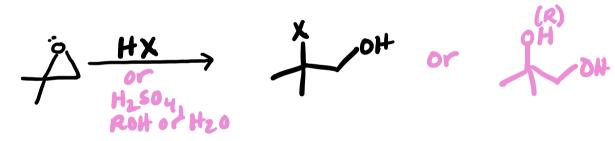
hork-no

- mechanism is SN2
- huck ophile attacks less substituted side of epox.
- side of epoxido (due to steries)
- two part rxn i) strong Nu ii) H30+ work-up
- :Na = any strong nucleophile (anything that can do SN2 and organom dallics) "H" as LIAIHY DH, OR, NH, NHR, NR2 CN, SH, SR, NJ, I, BY amines - moderat No, but Still Works

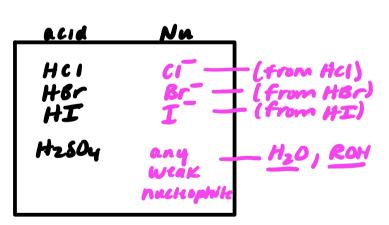
STEREOCHEMISTRY

SNZ - inversion of configuration at the carbon Nu attacks

Epoxide-Opening (acidic conditions)



- mechanism is SN2
- huck ophile attacks more substitued side of epox.
- side of epoxide (due to location of 5+ charge)
- One part run (strong acrd + weak Nu)



STEREOCHEMISTRY

SNZ - inversion of configuration at the carbon Nu attacks

H202

- oxidation to sufexide

Natoy

- oxidation to sulfone

- two SNZ, 2nd 15 intramolecular
- typical BN2 considerations apply