Chapter 13:Highlights

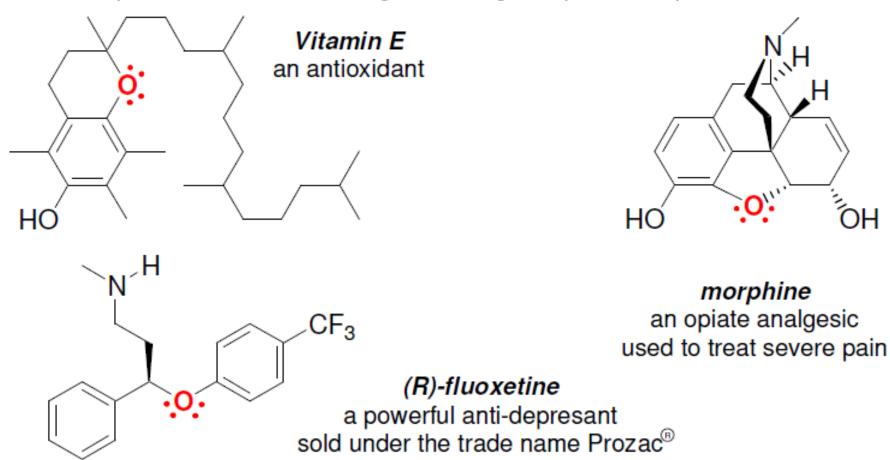
Today –(Ch. 13. 1- 13.6, 13.8)

Wednesday (Ch.13.9-13.12)

We will not cover nomenclature of epoxides.

CHAPTER 13: Introduction to Ethers

Compounds containing ether groups are quite common.

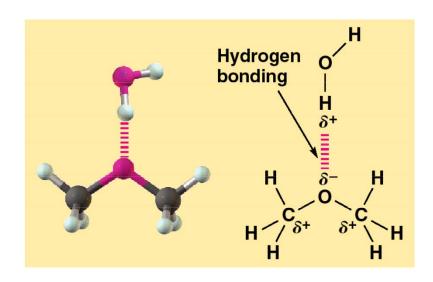


Ethers: Structure and Nomenclature

 An ether has two organic groups bonded to the same oxygen atom, R–O–R'

dimethyl ether





- Tetrahedral bond angle
- Oxygen is sp³-hybridized
- Oxygen atom gives ethers a slight dipole moment

Naming Ethers

For symmetrical ethers, name the alkane on each side and then add "ether."

For unsymmetrical (or asymmetrical) ethers, name the alkane on each side, and then add "ether."

CH₃CH₂OCH₂CH₃

CH₃CH₂CH₂CH₂O-

diethyl ether

Cyclohexyl butyl ether Butoxy-cyclohexane

As substituents, add "oxy" to the alkyl name:

Both are
Correct –
probably
see the
second
one more
often

CH₃O-

CH₃ CH₃CH₂O— CH₃

methoxy-

tert-butoxy-

If other functional groups are present, the ether part is considered an <u>alkoxy</u> substituent

Cyclic Ethers

$$H_2C$$
 CH_2
 H_2C
 CH_2
 CH_2

$$H_2C$$
 CH_2
 H_2C-CH_2

1,4-Dioxane

Tetrahydrofuran

Cyclic ethers behave like acyclic ethers, except if it is 3-membered

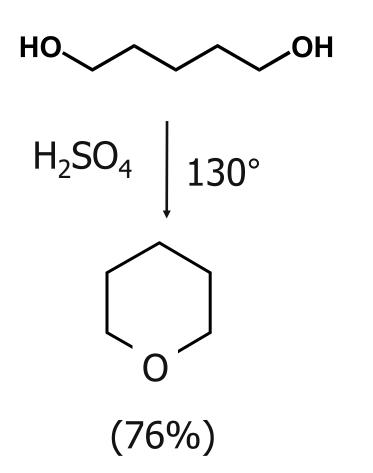


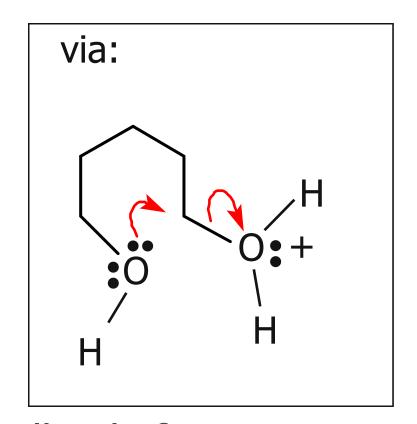
A three membered ring ether is called an oxirane or an EPOXIDE.

Synthesis of Ethers

Ethers are prepared industrially by acid—catalyzed dehydration of primary alcohols

Intramolecular Analog





reaction normally works well only for 5- and 6-membered rings

Preparation of Ethers

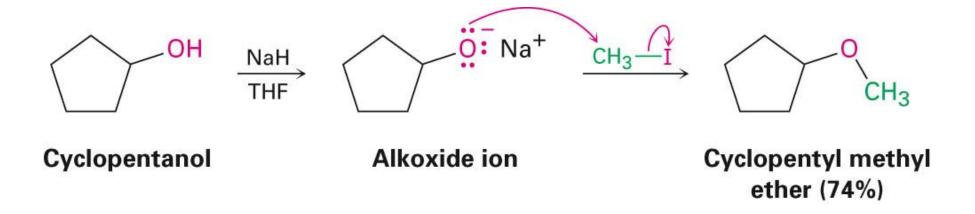
 Recall from Section 9.5 that oxymercurationdemercuration can be used to synthesize alcohols.

$$\begin{array}{c|c}
R & H \\
\hline
 & 1) Hg(OAc)_2, H_2O \\
\hline
 & 2) NaBH_4
\end{array}$$

 Similarly, alkoxymercuration-demercuration can be used to synthesize ethers.

The Williamson Ether Synthesis

- Reaction of metal alkoxides and primary alkyl halides and tosylates
- Best method for the preparation of ethers
- Alkoxides are prepared by reaction of an alcohol with a strong base Ex: NaH or NaNH₂



Preparation of Epoxides:

Treatment of an alkene with a peroxyacid

Treatment of a halohydrin with base gives an epoxide

Acid catalyzed Ring Opening of Epoxides

$$H_2^+$$
 H_2^+ H_2^+ $H_2^ H_2^ H_2^$

Nucleophile attacks <u>more</u> substituted carbon of protonated epoxide

Inversion of configuration at site of nucleophilic attack

Mild acid reacts with epoxides in water to generate anti 1,2-diols

Regioselectivity <u>under acidic conditions</u> explained if we consider that protonated epoxide has slight carbocation character.

$$\begin{array}{c} H \\ \delta^{+} \\ H_{3}CHC-CH_{2} \\ \hline \\ Br \\ \hline \\ resembles 1^{\circ} \\ carbocation \\ less stable \\ \hline \\ less product \\ \end{array}$$

REMEMBER: When a nucleophile attacks an <u>unprotonated</u> epoxide (under basic conditions) the reaction is a pure $S_N 2$ reaction

For Next Time....

Suggested Homework Problems

Chapter 13 #1, 5, 10, *18, 26, 29, 37, 38, 44, *51