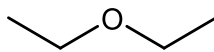
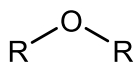


# Ethers



diethyl ether

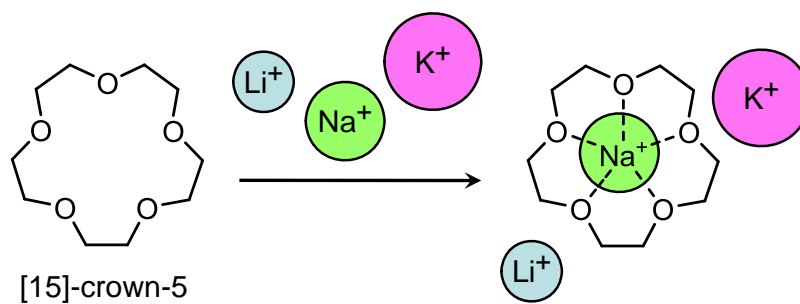
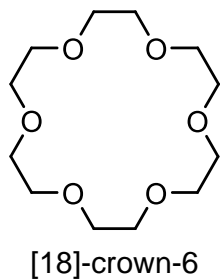
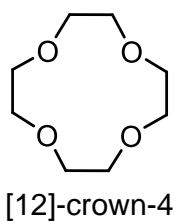


an epoxide  
(a cyclic ether)

Ethers are good Lewis bases/electron donors.

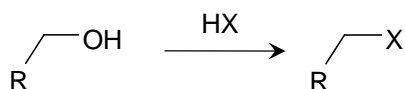
Example:  $\text{BH}_3 \cdot \text{THF}$  (hydroboration reagent):

# Crown Ethers

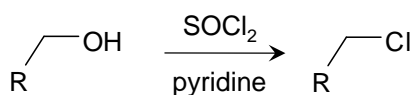
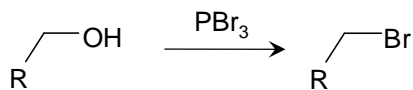


| ion           | ionic diameter | recognized by       |
|---------------|----------------|---------------------|
| $\text{Li}^+$ | 1.20 Å         | [12]-crown-4        |
| $\text{Na}^+$ | 1.80 Å         | <b>[15]-crown-5</b> |
| $\text{K}^+$  | 2.66 Å         | [18]-crown-6        |

## Direct Conversion of Alcohols to Halides



Not so great. E1/E2 compete with substitution.

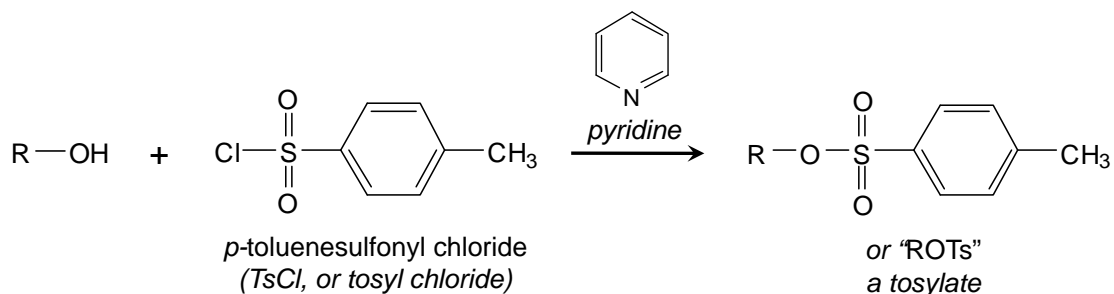


These reactions:

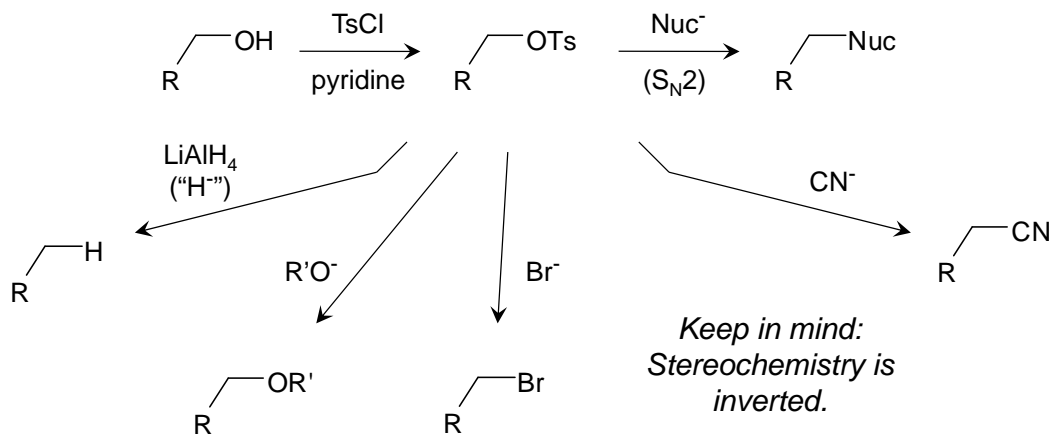
- convert  $-\text{OH}$  into a good leaving group
- invert stereochemistry at carbon.

*In the absence of pyridine, this reaction can retain stereochemistry.*

## Tosylation of Alcohols



## Tosylation Turns –OH Into an Excellent Leaving Group

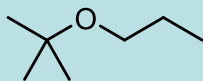


## Williamson Ether Synthesis

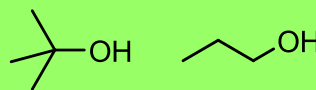
Ethers (though not epoxides) can be synthesized by reacting an alkoxide nucleophile with a good alkyl electrophile.

*Example*—From two alcohols:

How might we make this

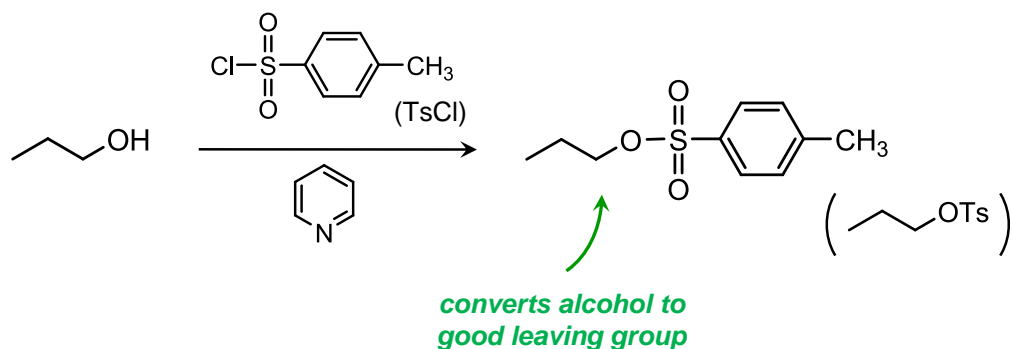


from these?



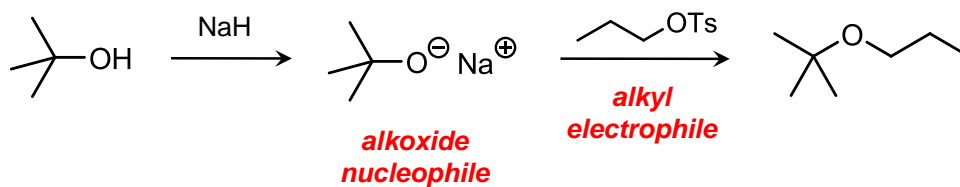
# Williamson Ether Synthesis

First, convert one alcohol to a good leaving group:



# Williamson Ether Synthesis

Second, combine tosylate with reactive alkoxide:



- Works well with a primary (1°) halide/tosylate;
- Not so well with 2° or 3° leaving groups; too much competing E2 elimination.