

Substitution Reactions – 2 types: S_N1 and S_N2

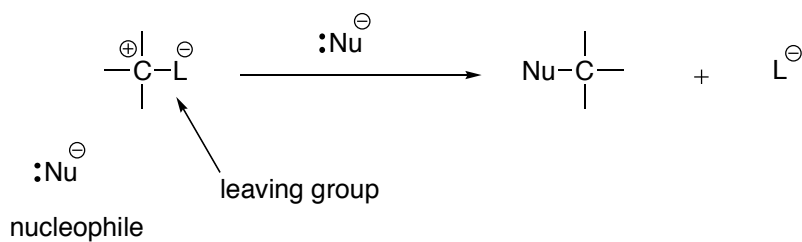
S_N2 reactions:

S = Substitution

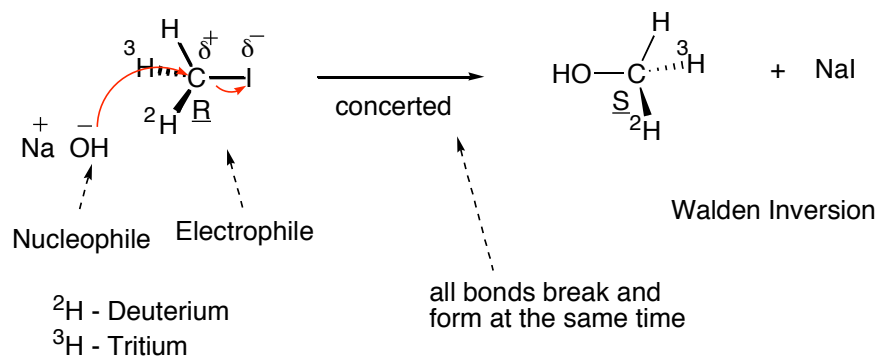
N = nucleophilic

2 = bimolecular reaction (rate of reaction depends on 2 reagents)

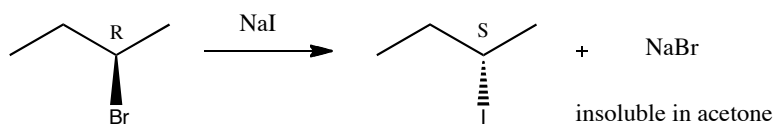
- stereospecific reaction
- inversion of configuration
- concerted reaction
- rate depends on two reagent concentration: [Nu] and [SM]
- favored for primary 1° carbons



Eg.

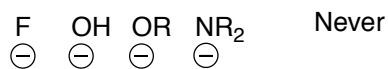


Eg.

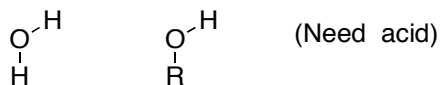


Leaving group ability

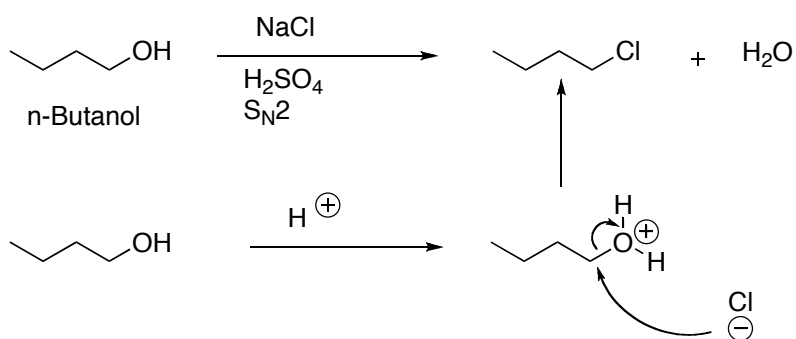
Worst



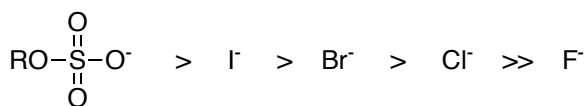
Possible Leaving Groups – ACID required



Eg.



Excellent to Good Leaving Groups

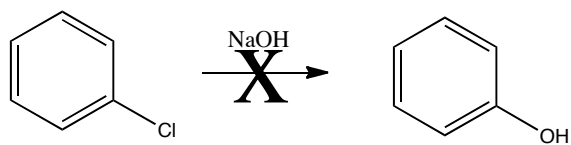


very good

poor

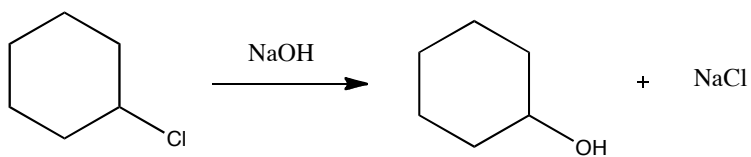
The order of halide leaving group ability is due to solvation and size.

Eg.

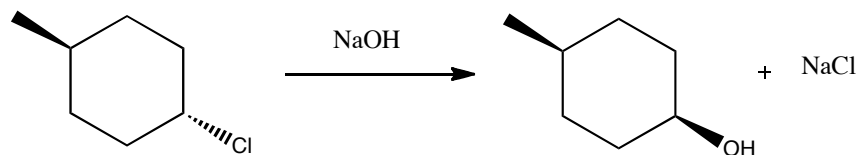


$\text{S}_{\text{N}}2$ **never** occurs at double bonds.

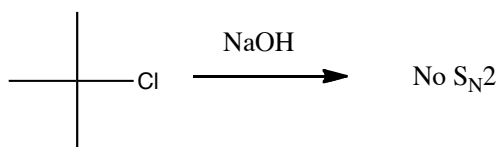
Eg.



Eg.



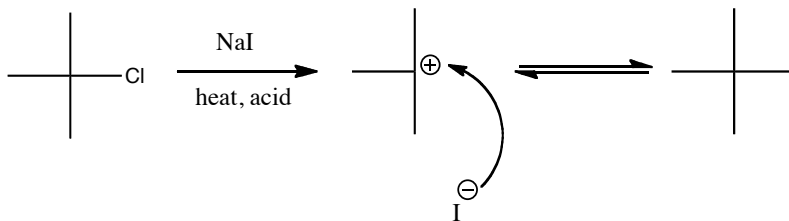
Eg.



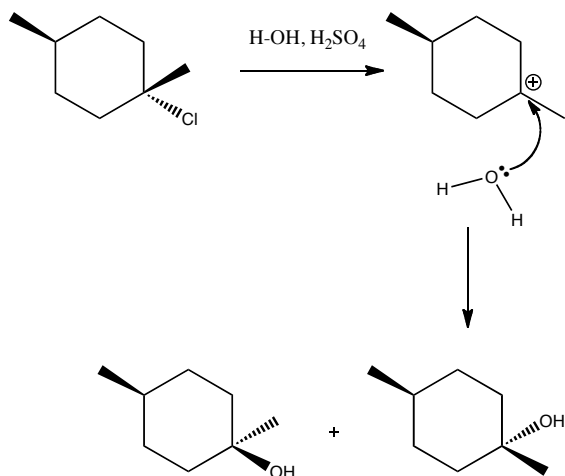
S_N1 reactions - Substitution Nucleophilic Unimolecular

- rate depends on 1 concentration: $[SM]$
- stepwise (not concerted)
- carbocation intermediate
- not stereospecific

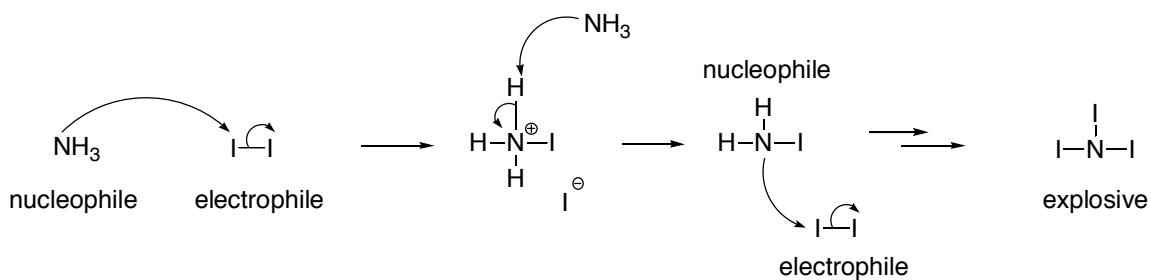
Eg.



Eg.



Demo:



S_N1

- Best if 3° carbocation can form
- Never on 1° alkyl halides
- Leaving groups – same as S_N2

OR and OH work if strong acid present
HOR and HOH are leaving groups

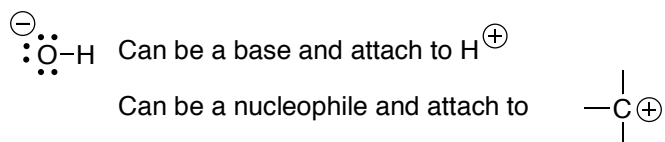
S_N1 and S_N2 can compete with E1 and E2 (alkenes formed)

Base : \ominus

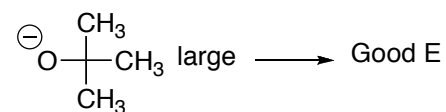
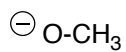
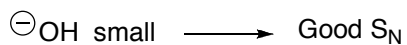
Nucleophile : \ominus

For elimination

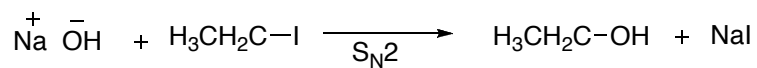
For substitution



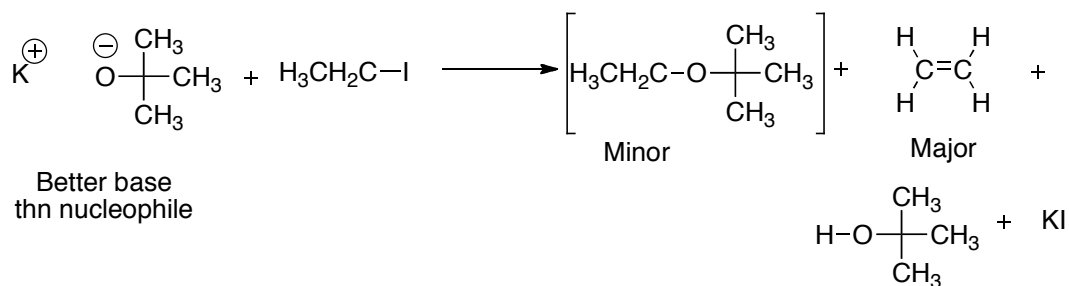
How can you tell which?



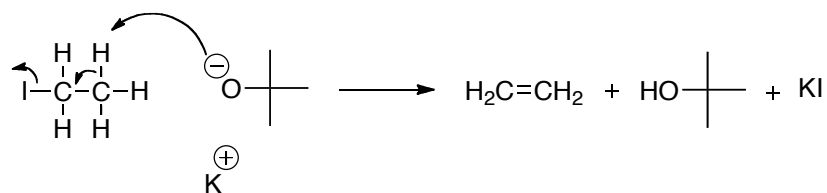
Eg.



Eg.

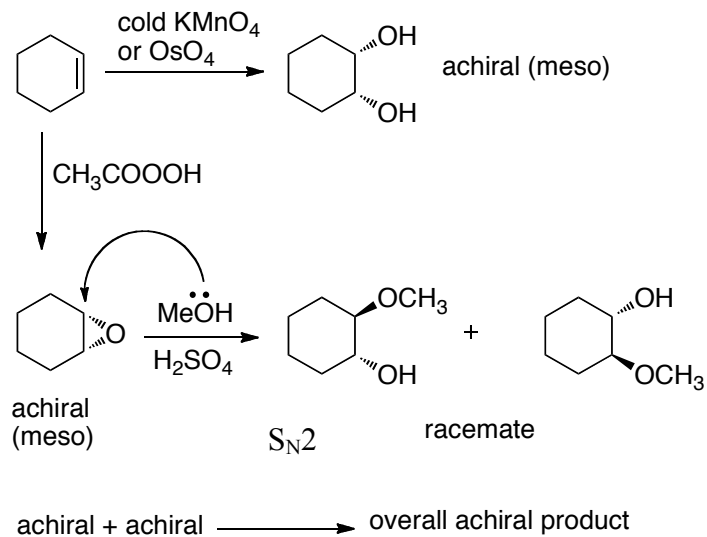


Eg.

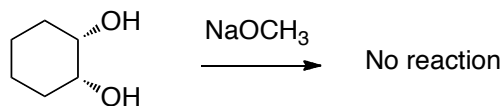


Review

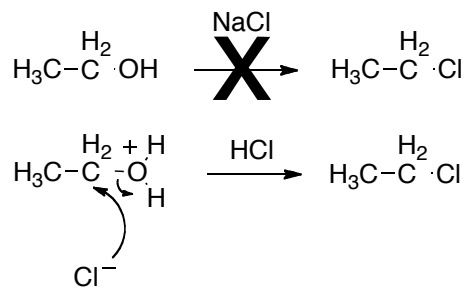
Eg.



Eg.



Eg.



Eg.

