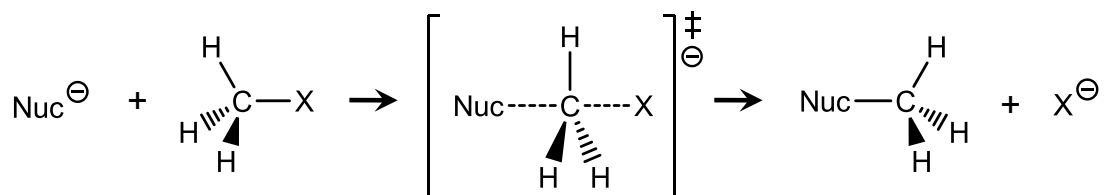


What Makes a Good Nucleophile?



1. Wants to give away electrons (good Lewis base).

HO^- better than H_2O

(bases always better than their conjugate acids)

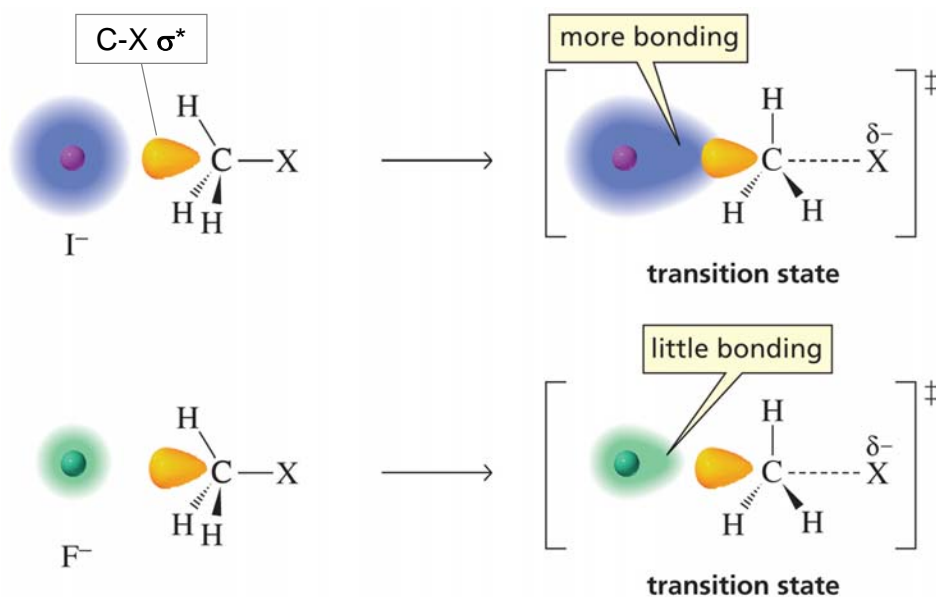
NH_3 better than H_2O

HS^- better than Cl^-

(less electronegative atom means more willing to give up e^-)

What Makes a Good Nucleophile?

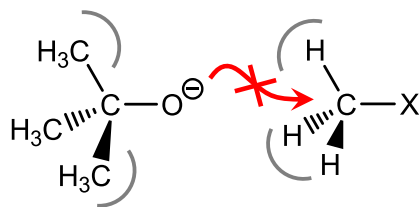
2. More polarizable. (Bigger.)



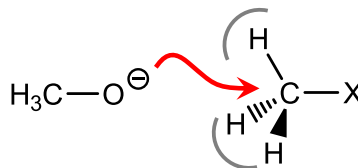
So, I^- is a better nucleophile than F^- .

What Makes a Good Nucleophile?

3. Not too sterically hindered.



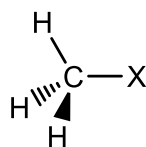
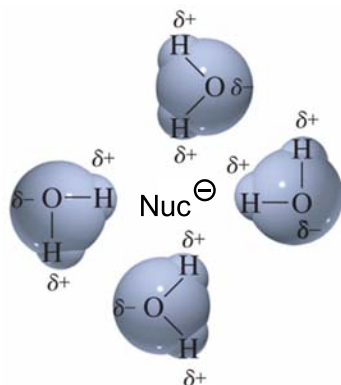
t-butoxide anion
strong base,
hindered nucleophile



methoxide anion
*smaller, attacks
more easily*

What Makes a Good Nucleophile?

4. Not obscured by too polar a solvent.



*solvent hinders
nucleophile from
approaching
electrophile.*

So, S_N2 reactions are usually run in medium polarity solvents;
polar enough to dissolve ionic species, not so polar to cage the nucleophile.

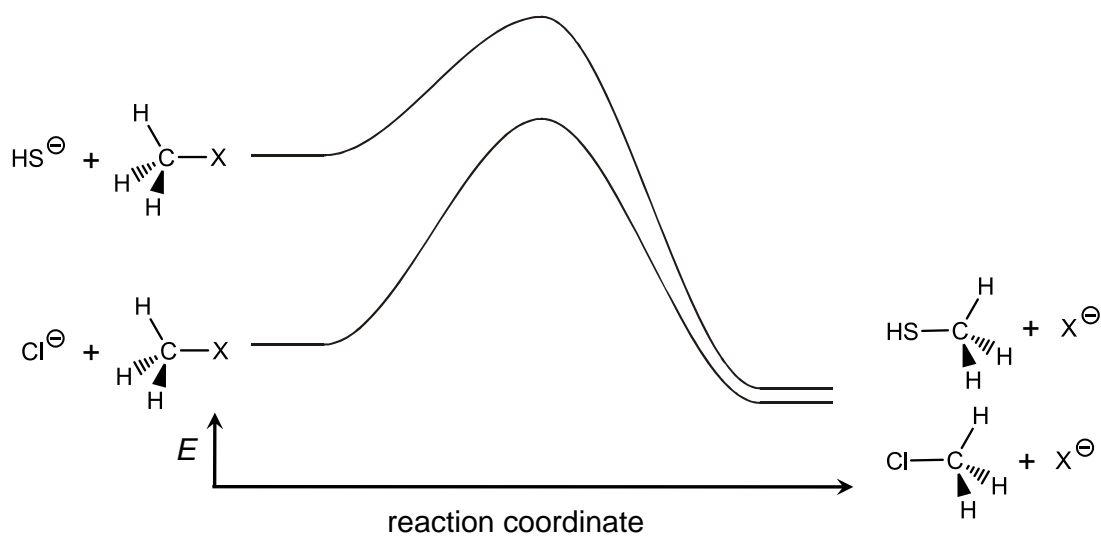
Some Common Nucleophiles

TABLE 6-3 Some Common Nucleophiles. Listed in Decreasing Order of Nucleophilicity in Hydroxylic Solvents Such as Water and Alcohols

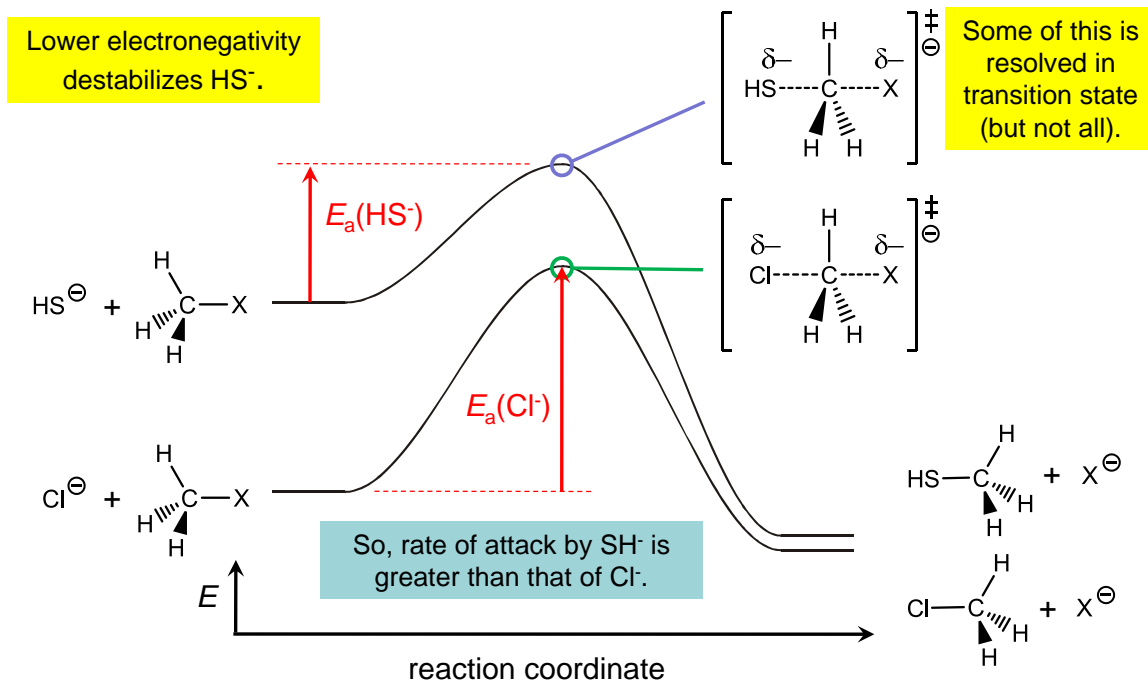
| | | | |
|------------------------------------|---|-----------------------|---|
| strong nucleophiles | $(\text{CH}_3\text{CH}_2)_3\text{P:}$ | moderate nucleophiles | $:\ddot{\text{Br}}:^{-}$ |
| | $^{-}:\ddot{\text{S}}-\text{H}$ | | $:\text{NH}_3$ |
| | $:\ddot{\text{I}}:^{-}$ | | $\text{CH}_3-\ddot{\text{S}}-\text{CH}_3$ |
| | $(\text{CH}_3\text{CH}_2)_2\ddot{\text{N}}\text{H}$ | | $:\ddot{\text{Cl}}:^{-}$ |
| | $^{-}:\text{C}\equiv\text{N}$ | | $\text{CH}_3\text{C}(=\text{O})-\ddot{\text{O}}:^{-}$ |
| | $(\text{CH}_3\text{CH}_2)_3\text{N:}$ | | $:\ddot{\text{F}}:^{-}$ |
| | $\text{H}-\ddot{\text{O}}:^{-}$ | | weak nucleophiles |
| $\text{CH}_3-\ddot{\text{O}}:^{-}$ | $\text{CH}_3-\ddot{\text{O}}-\text{H}$ | | |

Comparing Nucleophile Quality with Potential Energy Diagrams

We said HS^- better than Cl^- (based on electronegativity).

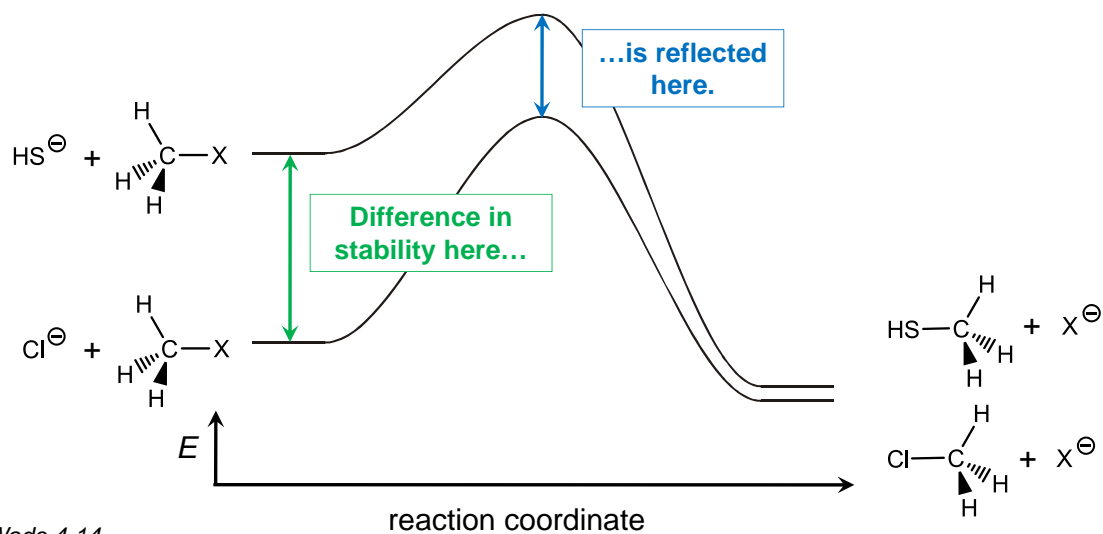


Comparing Nucleophile Quality with Potential Energy Diagrams



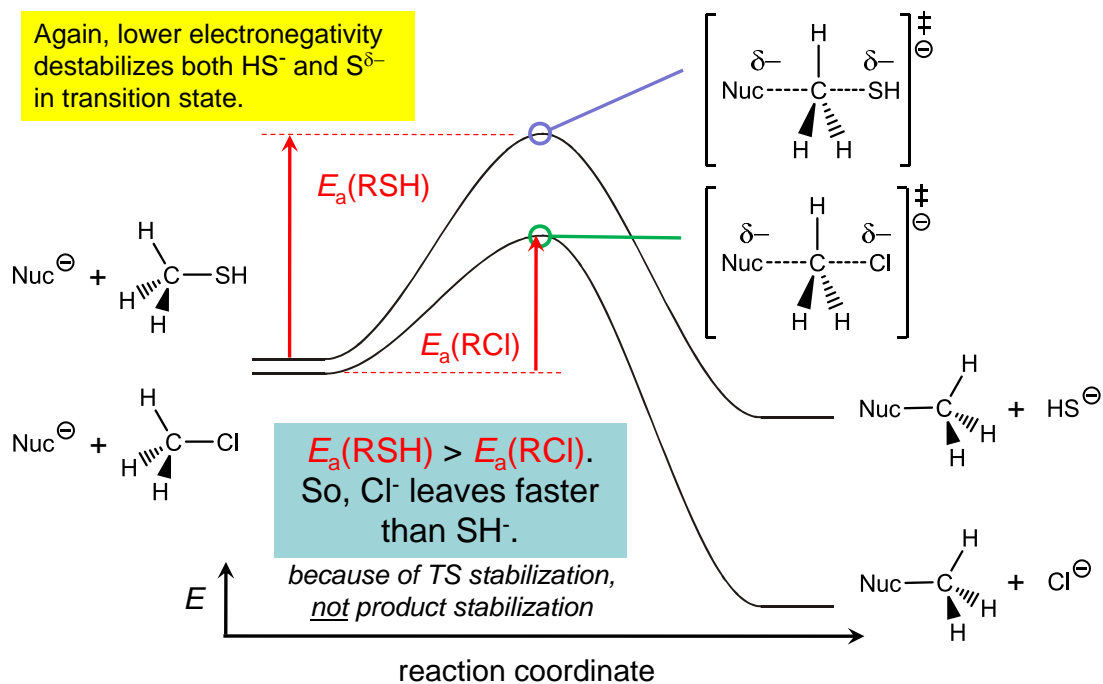
The Hammond Postulate

For similar reactions, differences in starting material or product stabilities will be reflected in transition state (to a lesser degree).



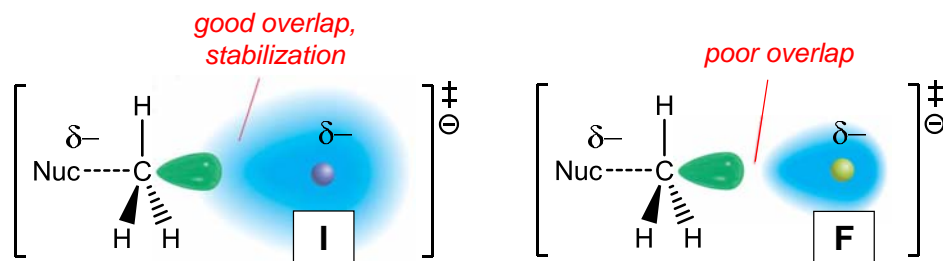
What Makes a Good Leaving Group?

1. Wants to take electrons (electronegative).



What Makes a Good Leaving Group?

2. Polarizable. (Stabilizes transition state.)



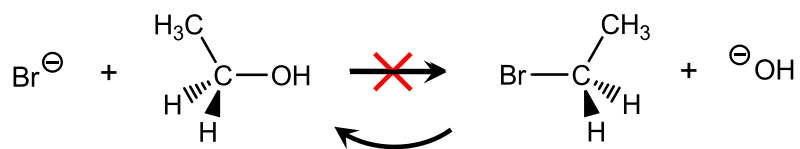
So, I^- is a better leaving group than F^- .

Interesting consequence: I^- is both a good nucleophile
and a good leaving group.

What Makes a Good Leaving Group?

3. Products are stable; S_N2 wouldn't work much better in reverse.

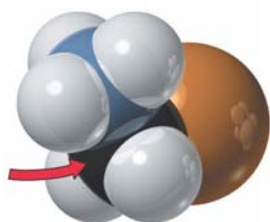
Example:



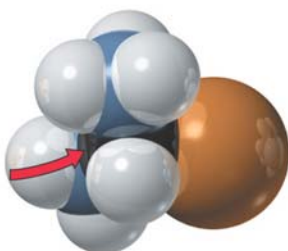
$\ominus\text{OH}$ is a much better nucleophile than Br^\ominus ; this reaction would revert if it ever happened.

So it doesn't happen.

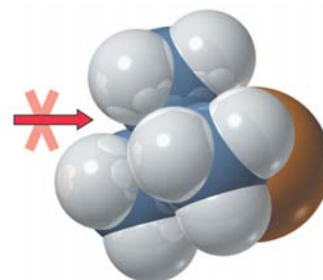
Steric Hindrance at the S_N2 Reaction Center Inhibits Reaction



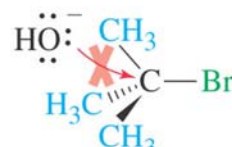
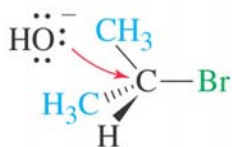
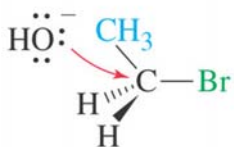
ethyl bromide (1°)
attack is easy



isopropyl bromide (2°)
attack is possible



t-butyl bromide (3°)
attack is impossible



Inhibition of S_N2 by Neopentyl Groups

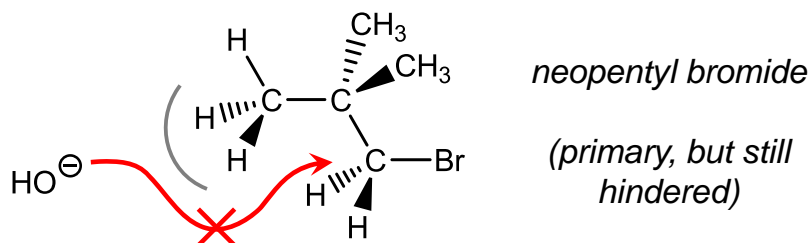


TABLE 6-5 Effect of Substituents on the Rates of S_N2 Reactions

| Class of Halide | Example | Relative Rate |
|----------------------|---|---------------|
| methyl | CH ₃ —Br | >1000 |
| primary (1°) | CH ₃ CH ₂ —Br | 50 |
| secondary (2°) | (CH ₃) ₂ CH—Br | 1 |
| tertiary (3°) | (CH ₃) ₃ C—Br | <0.001 |
| <i>n</i> -butyl (1°) | CH ₃ CH ₂ CH ₂ CH ₂ —Br | 20 |
| isobutyl (1°) | (CH ₃) ₂ CHCH ₂ —Br | 2 |
| neopentyl (1°) | (CH ₃) ₃ CCH ₂ —Br | 0.0005 |