Chapter 11 Nucleophilic Substitution $(S_N 1/S_N 2)$ Elimination (E1/E2)

Suggested Problems for Elimination:

11.14, 11.17-19, 11.20, 11.34, 11.40, 11.45-46, 11.49-50, 11.56, 11.59, 11.65, 11.68

Substitution Reactions ($S_N 2$ versus $S_N 1$)

S_N1

Mechanism:	Concerted	Two Steps (Look for carbocation rearrangements.)
Rate Equation:	Rate = k _r [R–X][Nuc]	Rate = k _r [R–X]
Stereochemistry:	Stereospecific (inversion)	Loss of Stereochemistry
Substrate:	Sterics: (methyl > 1° > 2°) No S _N 2 with 3°! Accesible to Nu: (benzylic > allylic > 1°)	Cation Stability (benzylic > allylic > 3° > 2°>>1°) No 1° or methyl R ⁺ without extra resonance stabilization!
Nucleophile:	Strong/Moderate Required strong: RS ⁻ , I ⁻ , R ₂ N ⁻ , R ₂ NH, RO ⁻ , CN ⁻ moderate: RSH, Br ⁻ , RCO ₂ ⁻	Not Important
Leaving Group:	Moderately Important (-OTf >> -OTs @ -OMs >> -I > -Br > -CI)	Very Important (-OTf >> -OTs @ -OMs >> -I > -Br > -CI)
Solvent:	Polar Aprotic	Polar Protic

Elimination Reactions: E2 versus E1

	E2	E1
Mechanism:	Concerted	Two Steps (Look for carbocation rearrangements.)
Rate Equation:	Rate = k _r [R–X][Base]	Rate = k _r [R–X]
Stereochemistry:	Stereospecific (antiperiplanar TS)	Not Stereospecific
Substrate:	Alkene Stability $(3^\circ > 2^\circ > 1^\circ)$	Cation Stability (benzylic > allylic > 3° > 2°>>1°)
Base:	Strong Base Required (RO ⁻ , R ₂ N ⁻)	Not Important: Usually Weak (ROH, R ₂ NH)
Leaving Group:	Moderately Important (same trend as S _N 1)	Very Important (same trend as S _N 1)
Solvent:	Wide Range of Solvents	Polar Protic
Product Patio:	Zaitsev Rule: The most highly substituted alkene usually predominates.	
	Hofmann Product: Use of a sterically hindered base will result in formation of the least substituted alkene (Hofmann product).	

Generic Reaction-Energy Diagrams





 * Under conditions that favor a unimolecular reaction (weak nuc/base and polar protic solvent), mixtures of S_N1 and E1 are usually obtained.