Course Outline

Nomenclature: Do it yourself. You are responsible for naming chains up to 12 carbons long for all classes of compound covered (alkanes, alkenes, alkynes, alcohols, thiols, halides, amines, aldehydes, ketones, ethers, cyclic and acyclic molecules). You are NOT required to name bicyclic compounds. A guide to nomenclature will be available on the slides presented in class when appropriate. In the text the basics of nomenclature are also covered. You should use the index to find rules for selected functional groups. Also as a carry-on from the last semester, refresh your memory with the following:

- Periodic table, electronegativity and periodic trends, covalent bonds
- Drawing organic structures: Lewis & compressed structures, formal charge method, polarity, resonance

Part A: Introduction
1) Organic Molecular structure (Ch 1-2)
   - Orbital shapes and hybridization. Bond angles and structure, σ and π bonds, geometries, relative energies
   - Dipoles, Van der Waals forces, H-bonding. Molecular structure and bulk properties
   - Drawing molecules
2) Alkanes (Ch 4)
   - Properties, line structures, σ bond rotation, conformations, energy potential diagrams, Newman projections
   - Cycloalkanes, ring strain, conformations, configurations, syn/anti and cis/trans nomenclature.
   - Chair conformation, axial and equatorial bonds, chair interconversions.
3) Stereochemistry (Ch 5)
   - Stereochemistry, configuration and R/S nomenclature, enantiomers, diastereomers and meso compounds
   - Zig-zag structures, syn/anti nomenclature, resolution, optical activity, enantiomeric excess.
4) Reaction mechanisms (Ch 6)
   - Refreshment of concepts of Thermo and Kinetics
   - Curved arrows use
   - Mechanisms and carbocation arrangements

Part B: Simple Reactivity
1) Acids and bases
   a. Acid/base reactions, acidity of organic compounds, inductive effects, resonance, periodic trends. Use of pKa values
   b. Functional groups, arrow notation, reaction co-ordinate diagrams.
2) π Bonds as electrophiles
b. Reduction of carbonyl groups. Additions “across” π bonds. Sodium borohydride and lithium aluminum hydride.
c. Imines. Acid catalysis.
d. Cyanohydrin formation. Reversal of the reaction. Acid and base catalysis
h. Acid and base catalysis in carbonyl additions. Oxonium ions. Additions to oxoniums.
i. Microscopic reversibility. Molecular orbitals and reactivity

**Part C: π Bonds and Reactivity**

1) π Bonds as nucleophiles

2) Aromatic compounds as nucleophiles
   b. Allyl groups and resonance. Allyl cation and anion.