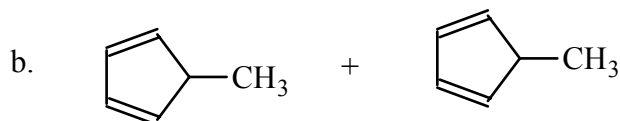
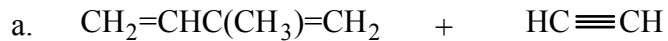


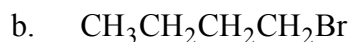
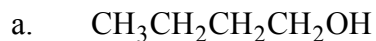
1. Give the products of the following Diels-Alder reactions. (10 pts)



2. What is the major product formed when trans-1-bromo-2-ethylcyclohexane undergoes an E2 elimination? Why? (5 pts)

3. Would it be better to use ^1H NMR or ^{13}C NMR to distinguish among 1-butene, cis-2-butene, and 2-methylpropene? Explain your answer.

4. Show, using any necessary reagents, how the following compounds could be prepared using ethylene oxide as one of the reactants:



5. What carbon is protonated in the first step of the reaction of 3-methyl-2,4-heptadiene with one equiv of HBr?

- a. C-2
- b. C-1
- c. C-3
- d. C-4
- e. C-5

6. Why do many $\text{S}_{\text{N}}1$ reactions produce more of the product with inverted configuration than with retained configuration?

- a. Reaction occurs by way of a Walden inversion. The product with retained configuration comes from racemization after the $\text{S}_{\text{N}}1$ reaction.
- b. The leaving group partially blocks the side of the carbocation from which it leaves.
- c. The solvent-separated ion pair results in more of the product with retained configuration.
- d. The nucleophile attacks the free carbocation.

7. Which of the following statements describes the reaction coordinate diagram for an E_1 reaction?

- a. a one step reaction with one transition state
- b. a two step reaction with 1 intermediate and 1 transition state
- c. a one step reaction with 2 transition states
- d. a one step reaction without a transition state
- e. a two step reaction with 1 intermediate and 2 transition states

8. Why does cis-1-bromo-4-tert-butylcyclohexane react faster in an E_2 reaction than does trans-1-bromo-4-tert-butylcyclohexane?

- a. Since the Br is in an equatorial position, syn elimination can occur.
- b. When Br and H are in the required equatorial positions, the large tert-butyl group is in the more stable axial position.

- c. The required syn elimination can occur with all the substituents in the most stable equatorial position.
- d. When Br and H are in the required axial positions, the large tert-butyl group is in the more stable equatorial position.
- e. When Br and H are in the required equatorial positions, the large tert-butyl groups is in the more stable equatorial position.

9. Which of the following best describes the carbon-lithium bond?

- a. The carbon-lithium bond is covalent and somewhat polar.
- b. The carbon-lithium bond is covalent and nonpolar.
- c. The carbon-lithium bond is almost ionic with carbon negative and lithium positive.
- d. The carbon-lithium bond is almost ionic with carbon positive and lithium negative.
- e. A carbon-lithium bond is less polar than a carbon-oxygen bond.

10. Which of the following is **not** true about electromagnetic radiation?

- a. Energy is directly proportional to the velocity of light.
- b. Frequency is directly proportional to wavelength.
- c. Frequency is directly proportional to wavenumber.
- d. Energy is inversely proportional to wavelength.
- e. Energy is directly proportional to wavenumber.

11. The ^1H NMR spectrum of a compound shows methyl and methylene signals with relative intensities of 3:1. Identify the compound.

- $\text{CH}_3\text{CH}_2\text{Br}$
- $(\text{CH}_3)_3\text{CCH}_2\text{Br}$
- $(\text{CH}_3)_2\text{CBrCH}_2\text{Br}$
- $(\text{CH}_3)_2\text{C}(\text{CH}_2\text{Br})_2$
- $\text{CH}_3\text{C}(\text{CH}_2\text{Br})_3$

12. How many peaks would be observed in the ^1H and ^{13}C NMR spectra respectively of 4-bromotoluene?

- 7 and 7
- 2 and 2
- 5 and 5
- 3 and 5
- 4 and 7

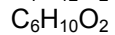
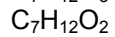
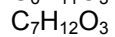
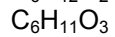
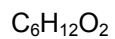
13. What is "ferrocene"?

- a complex of Fe^{3+} and 1 cyclopentadienyl dianion
- a complex of Fe^{2+} and 2 benzene anions
- a complex of Fe^{3+} and 2 cyclopentadienyl anions
- a complex of Fe^{2+} and 1 cyclopentadienyl dianion
- a complex of Fe^{2+} and 2 cyclopentadienyl anions

14. Which of the following compounds cannot be converted to benzoic acid by heating with an acidic solution of sodium dichromate?

- 2-methyl-2-phenylbutane
- 2-methyl-3-phenylbutane
- ethylbenzene
- 2-phenylbutane
- 1-phenylpropane

15. What is the molecular formula of 3-methylcyclopentanecarboxylic acid?



16. Draw the structures of the following molecules:

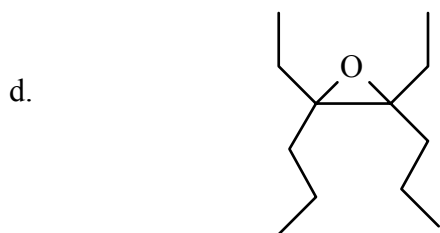
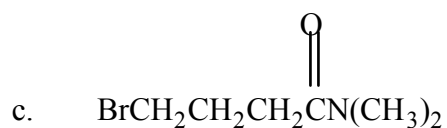
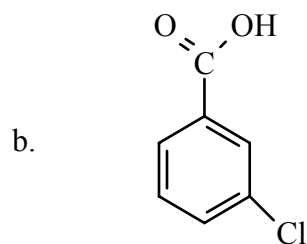
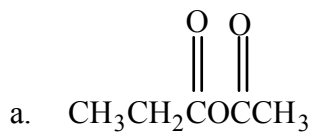
a. 6-methyl-2-cyclohexenol

b. 2-propyloxirane

c. 5-bromo-2-nitrotoluene

d. methyl 3-bromobutanoate

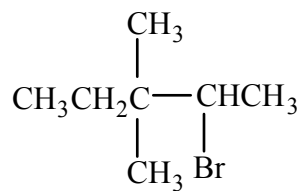
17. Name the following compounds:



18. Give the major 1,2- and 1,4-addition products for the following reaction, and indicate which is the thermodynamic and which is the kinetic product.



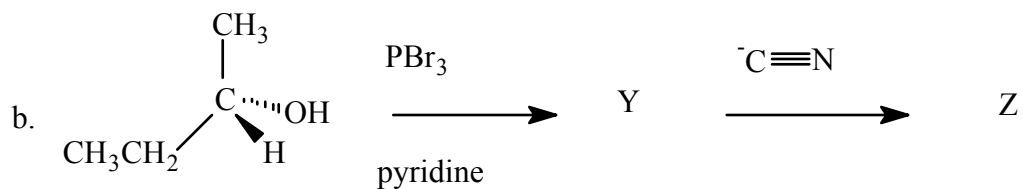
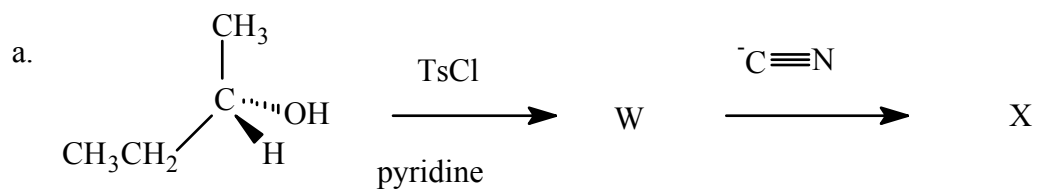
19. The following alkyl halide forms a substitution product from an $\text{S}_{\text{N}}1$ reaction that is different from the substitution product formed from an $\text{S}_{\text{N}}2$ reaction. Show the alcohol products formed from an $\text{S}_{\text{N}}1$ and an $\text{S}_{\text{N}}2$ reaction.



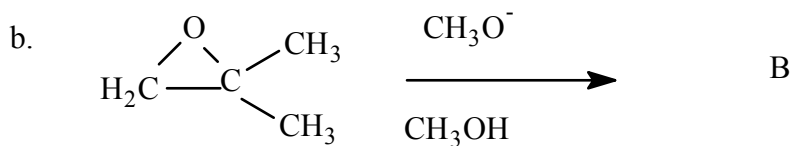
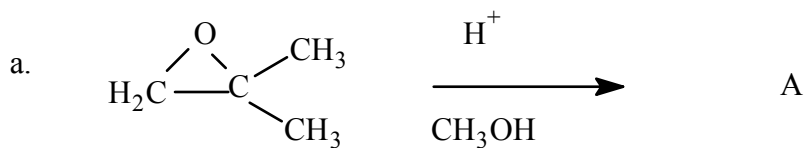
20. What alkene will be formed in an E2 reaction of the following compound. Indicate its configuration.

(1S, 2S)-1-bromo-1,2-diphenylpropane

21. Give the structures of W, X, Y, and Z.



22. Give the structures of A and B.



23. The mass spectra of two different cycloalkanes both show a molecular ion peak at $m/z = 98$. One spectrum shows a base peak at $m/z = 69$, and the other shows a base peak at $m/z = 83$. Identify the cycloalkanes. Explain!

24. Draw the π molecular orbital energy levels for the cyclopentadienyl anion, the cyclopentadienyl cation, and the cyclobutadiene. For each compound, show the distribution of the π electrons. Which of the compounds are aromatic? Which are antiaromatic?

25. Show how 1-bromo-2-phenylethane can be prepared from benzene.

26. Show how m-bromopropylbenzene can be prepared from benzene.

27. Write the mechanism for the reaction of acetyl chloride with water to form acetic acid.