Alt. Exam 1

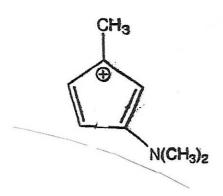
1. Give an acceptable IUPAC name for each of the compounds in a-c. Draw the structure of the compound in d. Be sure to indicate the stereochemistry where appropriate. (16 points)

b.

C.

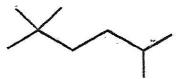
| 2. Draw a proper Lewis s<br>all lone pairs and formal                     | structure for the condense<br>charges. (5pts) | ed formula below. Be s                          | ure to show  |
|---|---|---|--------------|
|   |   |   |              |
| (CH <sub>3</sub> ) <sub>2</sub> OAICI <sub>3</sub>                        |   |   |              |
|   |   |   |              |
|   |   |   |              |
| <ol> <li>Place the compounds</li> <li>3=highest melting point)</li> </ol> | below in order of increas<br>(6 pts.)         | ing meiting point, (1=lo                        | owest,       |
|   |   |   | each.        |
|   |   |   |              |
|   |   |   |              |
| 4. Place the compounds soluble, 3=most soluble)                           | below in order of increas<br>(6 pts.)         | ing solubility in H <sub>2</sub> O. (1          | l=least      |
|   |   |   | 2pts<br>each |
| CH <sub>3</sub> CH <sub>2</sub> OCH <sub>2</sub> CH <sub>3</sub>          | CH <sub>3</sub> CH <sub>2</sub> OH            | CH <sub>3</sub> CH <sub>2</sub> NH <sub>2</sub> |              |
|   |   |   |              |

5. For the structure given below, draw the important resonance contributors. Circle the major contributor. (12 pts.)

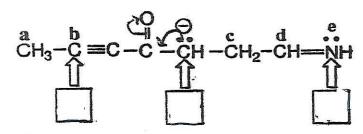


6. Place the compounds in order of increasing acidity. (1=least acidic, 3=most acidic) (6 pts.)

7. Draw all structural isomers resulting from the monobromination of 2,2,5-trimethyhexane (shown below). You will be penalized for duplicate structures. (10 pts.)



8. Consider the structure below and answer the following questions.



- a. Write the hybridization of each atom indicated by an arrow in the box provided. (6 pts.)
- b. What is the  $C_c$ - $C_d$ - $N_e$  bond angle? (2 pts.)
- c. The sigma bond between the atoms labeled a and b is formed by the overlap of what types of orbitals? Be specific. (2 pts.)

9. Viewing the molecule along the C3-C4 bond, construct the Newman projection of the least stable conformation of 3,4-dimethylheptane. (5 points)



10. a. Draw the more stable chair conformation for each of the substituted cyclohexanes shown below. (8 points)

b. Which isomer is more stable? A or B? (2 points)

| 44  | An equation representing the halogenation of an alkane is shown below.  | a) Provide the |
|-----|---|----------------|
| 11. | All equation representing the randy                                     | ate (6 naints) |
| ord | pagation steps of the mechanism that explain the formation of the produ | cis. (a house) |

$$+$$
 Br<sub>2</sub>  $\xrightarrow{\text{light}}$   $+$  HBr

- b. Consider the termination steps, and provide a structure for a possible side product. (2pt.)
- c. Given the bond dissociation energies (BDE) below, calculate the overall  $\Delta$  H°for the reaction, a You must show your work to receive credit! (4 points)
- d. Are the products or reactants favored at equilibrium? (2 points)

| Bond-Dissociation<br>Energy   |   |   | Bond-Dissociation<br>Energy  |  |
|---|---|---|--|--|
| Bond  | kcal/mol  | Bond  | kcal/mol :   |  |
| H—X bonds and X—X bonds H—H D—D F—F Cl—Cl Br—Br I—I H—F H—Cl H—Br H—I HO—H HO—OH Methyl bonds CH <sub>2</sub> —H CH <sub>3</sub> —F CH <sub>3</sub> —F CH <sub>3</sub> —G CH <sub>3</sub> —Br CH <sub>3</sub> —I CH <sub>3</sub> —OH Bonds to primary carbons CH <sub>3</sub> CH <sub>2</sub> —H CH <sub>3</sub> CH <sub>2</sub> —H CH <sub>3</sub> CH <sub>2</sub> —H CH <sub>3</sub> CH <sub>2</sub> —F CH <sub>3</sub> CH <sub>2</sub> —H CH <sub>3</sub> CH <sub>2</sub> —F CH <sub>3</sub> CH <sub>2</sub> —Br CH <sub>3</sub> CH <sub>2</sub> —I CH <sub>3</sub> CH <sub>2</sub> —Cl CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> —Br | 104<br>106<br>38<br>58<br>46<br>36<br>136<br>103<br>88<br>71<br>119<br>51<br>104<br>109<br>84<br>70<br>56<br>91<br>98<br>107<br>81<br>68<br>53<br>91<br>98<br>107<br>81<br>68<br>53 | Bonds to secondary carbons  (CH <sub>3</sub> ) <sub>2</sub> CH—H  (CH <sub>3</sub> ) <sub>2</sub> CH—F  (CH <sub>3</sub> ) <sub>2</sub> CH—E  (CH <sub>3</sub> ) <sub>2</sub> CH—B <sub>1</sub> (CH <sub>3</sub> ) <sub>2</sub> CH—I  (CH <sub>3</sub> ) <sub>2</sub> CH—I  (CH <sub>3</sub> ) <sub>2</sub> CH—I  (CH <sub>3</sub> ) <sub>3</sub> C—H  (CH <sub>3</sub> ) <sub>3</sub> C—H  (CH <sub>3</sub> ) <sub>3</sub> C—F  (CH <sub>3</sub> ) <sub>3</sub> C—F  (CH <sub>3</sub> ) <sub>3</sub> C—C  (CH <sub>3</sub> ) <sub>3</sub> C—C  (CH <sub>3</sub> ) <sub>3</sub> C—OH  Other C—H bonds  PhCH <sub>2</sub> —H (benzylic)  CH <sub>2</sub> —CHCH <sub>2</sub> —H (allylic)  CH <sub>2</sub> —CH—H (vinyl)  Ph—H (aromatic)  C—C bonds  CH <sub>3</sub> —CH <sub>3</sub> CH <sub>3</sub> —CH <sub>3</sub> CH <sub>3</sub> CH <sub>2</sub> —CH <sub>3</sub> CH <sub>3</sub> CH <sub>2</sub> —CH <sub>3</sub> (CH <sub>3</sub> ) <sub>3</sub> C—CH <sub>3</sub> | 95<br>106<br>80<br>68<br>53<br>91<br>91<br>106<br>79<br>65<br>50<br>91<br>85<br>87<br>108<br>110 |  |