NAME (Print):	·	 Dr.	emistry 310N Brent Iverson
SIGNATURE:			t Midterm b. 19, 2009
	Please print the first three letters of your last name in the three boxes		

**Please Note:** This test may be a bit long, but there is a reason. I would like to give you a lot of little questions, so you can find ones you can answer and show me what you know, rather than just a few questions that may be testing the one thing you forgot. **I recommend you look the exam over and answer the questions you are sure of first**, then go back and try to figure out the rest. Also make sure to **look at the point totals** on the questions as a guide to help budget your time.

For synthesis problems GO FOR PARTIAL CREDIT EVEN IF YOU DO NOT KNOW THE ENTIRE ANSWER!!!WRITE DOWN WHAT YOU DO KNOW IS IN THE REACTION SEQUENCE SOMEWHERE. YOU WILL GET PARTIAL CREDIT IF IT IS CORRECT

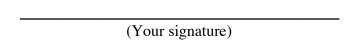
Note: You must have your answers written in pen if you want a regrade!!!!

Page	Points	•
1		(23)
2		(11)
3		(26)
4		(6)
5		(5)
6		(5)
7		(5)
8		(5)
9		(-)
10		(5)
11		(12)
12		(14)
13		(35)
14		(23)
15		(18)
16		(10)
17		(13)
18		(13)
19		(19)
Total		(248)
%		
T Score		
HW		
Total Grade		

(HW score + Exam Grade)

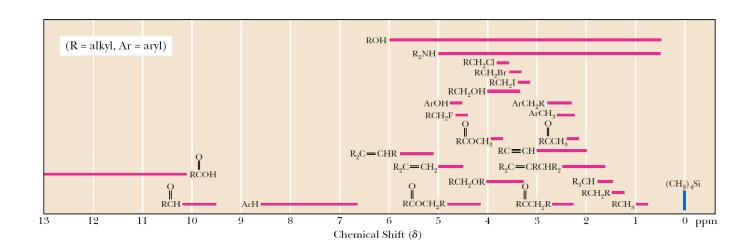
## **Honor Code**

The core values of the University of Texas at Austin are learning, discovery, freedom,
leadership, individual opportunity, and responsibility. Each member of the University is
expected to uphold these values through integrity, honesty, trust, fairness, and respect
toward peers and community.



Type of Hydrogen (R = alkyl, Ar = aryl)	Chemical Shift (δ)*	Type of Hydrogen (R = alkyl, Ar = aryl)	Chemical Shift (δ)*
	51111 (5)	(1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	51111 (0)
		RCH <sub>2</sub> OH	3.4-4.0
R <sub>2</sub> N <b>H</b>	0.5-5.0	RCH₂Br	3.4-3.6
ROH	0.5-6.0	RCH <sub>2</sub> Cl	3.6-3.8
RCH <sub>3</sub>	0.8-1.0	0	3.0 3.0
RCH <sub>2</sub> R	1.2-1.4	RCOCH3	3.7-3.9
R <sub>3</sub> C <b>H</b>	1.4-1.7	0	
R <sub>2</sub> C=CRCHR <sub>2</sub>	1.6-2.6	RCOCH2R	4.1-4.7
RC <b>≡</b> C <b>H</b>	2.0-3.0	RCH <sub>2</sub> F	4.4-4.5
O .		ArOH	4.5-4.7
RCCH <sub>3</sub>	2.1-2.3	$R_2C=CH_2$	4.6-5.0
0		R₂C=C <b>H</b> R	5.0-5.7
RCCH2R	2.2-2.6	0	
ArC <b>H</b> <sub>3</sub>	2.2-2.5	H <sub>2</sub> G—CH <sub>2</sub>	3.3-4.0
RCH <sub>2</sub> NR <sub>2</sub>	2.3-2.8	- <b>Ⅱ</b>	
RCH <sub>2</sub> I	3.1-3.3	R <b>CH</b>	9.5-10.1
RCH <sub>2</sub> OR	3.3-4.0	RCOH	10-13

<sup>\*</sup> Values are relative to tetramethylsilane. Other atoms within the molecule may cause the signal to appear outside these ranges.



- 1. (3 pts) For this class, the most important question in organic chemistry is:
- **2.** (14 points) Suppose a relative of yours is having an MRI. In no more than four sentences, explain to them what is happening when they have the MRI scan. We will be looking for a minumum of 7 key points here.

**3.** (3 pts each) Write an acceptable IUPAC name for the following molecules:

A.

$$\begin{array}{c|c} CH_3 & O \\ | & | \\ H_3C-C-CH_2-C-H \\ | \\ CH_3 \end{array}$$

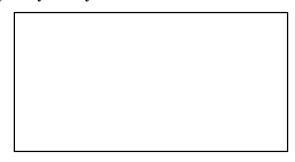
В.

Signature		
Jigiiatai C		

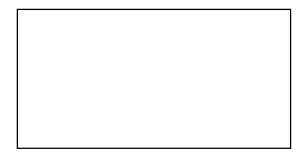
$P_{\sigma}$	2	(	11	)
15	_		11	,

**4.** (3 pts each) Write a structure that corresponds to the following IUPAC names:

A. (2S, 3S)-3-Hydroxy-2-ethylhexanal



B. 3-Methyl-3-butene-2-one



5. (1 pt each) Fill in each blank with the word that best completes the following sentences about NMR.

In the NMR experiments most useful to organic chemists, the nuclei \_\_\_\_\_\_ and \_\_\_\_\_ are the most important since these are commonly found in organic molecules and they have spin quantum numbers of 1/2. Nuclei with spin quantum number 1/2 are quantized in one of two orientations, "\_\_\_\_\_ " (lower energy) or "\_\_\_\_\_ "(higher energy) in the presence of an external magnetic field, that is, with and against the external field, respectively. The difference in energy between nuclear spin states is \_\_\_\_\_\_ to the strength of the magnetic field experienced by the nucleus.

Pg 3 \_\_\_\_\_(26)

**6.** (2 pts each) **Circle** all of the following statements about NMR that are **true** (do nothing if the statements are false).

If electromagnetic energy of the precise energy difference between the +1/2 and -1/2 spins states is used to irradiate the sample, the energy is absorbed as the spin "flips" from the +1/2 state to the -1/2 state.

If electromagnetic energy of the precise energy difference between the +1/2 and -1/2 spins states is used to irradiate the sample, the energy is absorbed as the spin "flips" from the -1/2 state to the +1/2 state.

"Resonance" in NMR refers to the phenomenon of absorption of energy when a nuclear spin "flips."

Equivalent hydrogen atoms in a molecule give the same NMR signal.

Equivalent hydrogen atoms in a molecule have identical relationships to all the other atoms in the molecule.

Hydrogen atoms in a molecule are equivalent if they are attached to the same freely rotating  $sp^3$  hybridized carbon atom.

Hydrogen atoms in a molecule are equivalent if they are in groups that are related by a symmetry element, such as being reflections of each other when the molecule is bisected by a plane of symmetry.

Two hydrogen atoms in a molecule are equivalent if they are attached to a carbon atom that is part of a non-symmetrical three-membered ring.

The physical basis of spin-spin splitting is that nuclei no more than three bonds away from the hydrogen of interest have a small magnetic field associated with their +1/2 or -1/2 spin state.

The physical basis of shielding is that electron density is induced to circulate in a strong magnetic field, and this circulation produces its own magnetic field that opposes the external magnetic field.

The physical basis of spin-spin splitting is that electron density is induced to circulate in a strong magnetic field, and this circulation produces its own magnetic field that opposes the external magnetic field.

THEORY: The signal for a  $-CH_2$  - group with a  $-CH_2$  - group and a  $-CH_3$  group on either side could show up to 3 x 4 = 12 peaks.

PRACTICE: For alkyl groups with freely rotating  $sp^3$  hybridized carbon atoms, complex splittings simplify because coupling constants ("J") are all about the same. In practice, if there are n adjacent H atoms, equivalent or not, you will see n+1 peaks. This is an approximation, but almost always true on spectra taken with all but the most sophisticated NMR spectrometers.

7. (1 pt each) Fill in each blank with the word that best completes the following sentences about NMR.

In the so-called "FT" approach to NMR, al	ll the nuclearare flipped
instantaneously with a multi-frequency pu	lse, then the rate at which the
"relax" back to the	state is monitored. This latter approach uses
a mathematical algorithm called	
(FT) to reconstruct individual resonance	for the different
equivalent sets of hydrogen atoms so the sp	ectra can be plotted.

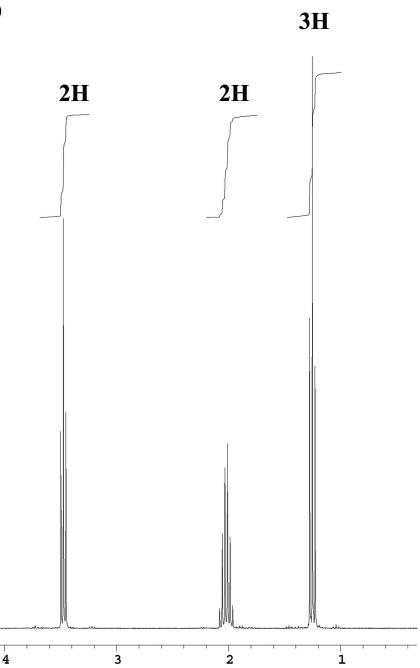
Here are some recently discovered molecular insects.

ppm

**8**. (5 pts each) Circle the molecule that corresponds to the spectrum shown.

$$\longrightarrow$$
 Br  $\longrightarrow$  OH  $\longrightarrow$  NH<sub>2</sub>

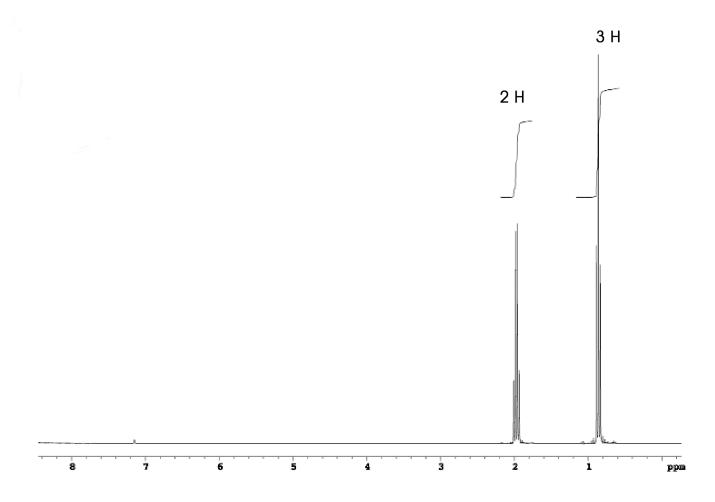
(Note: No  $D_2O$  has been added)



**9.** (5 pts) Circle the structure that corresponds to the following spectrum.

$$\bigcirc O \longrightarrow O \longrightarrow O \longrightarrow H$$

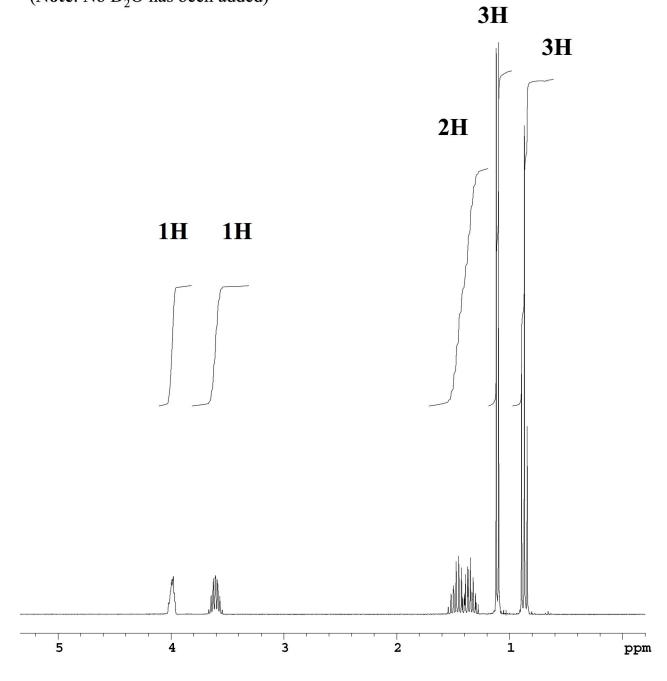
(No D<sub>2</sub>O was added to the sample)



10. (5 pts each) Circle the molecule that corresponds to the spectrum shown.

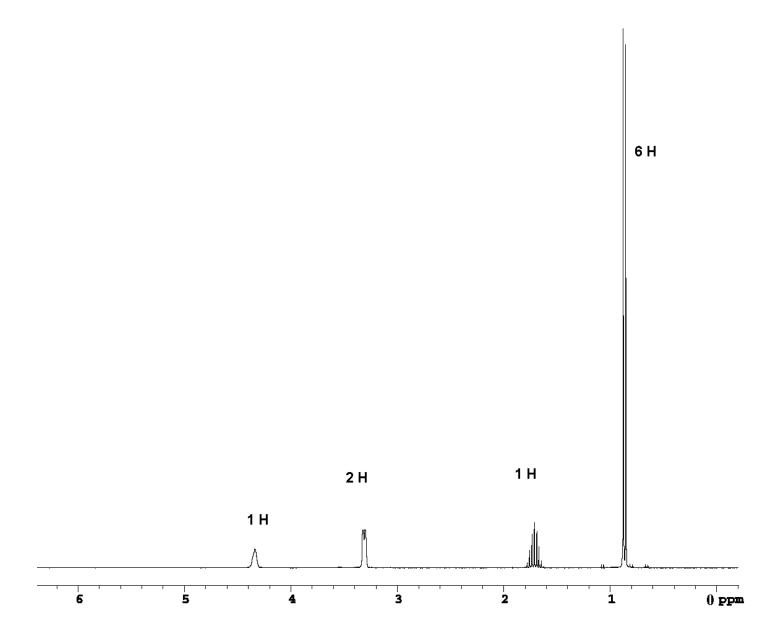
$$OH$$
  $OH$   $OH$   $OH$ 

(Note: No  $D_2O$  has been added)



11. (5 pts) Circle the structure that corresponds to the following spectrum.

(No  $\mathrm{D}_2\mathrm{O}$  was added to the sample)



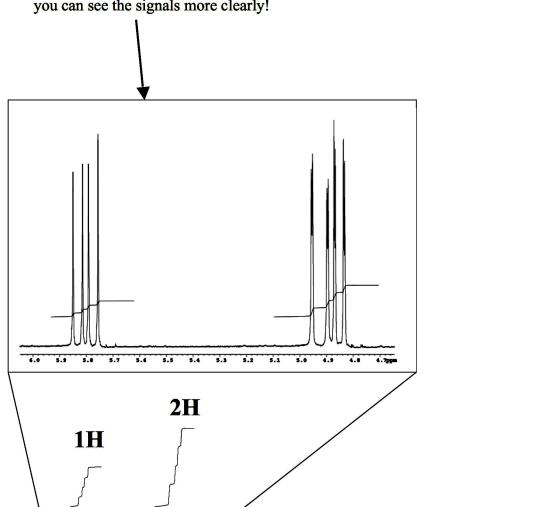
13. (5 pts each) Circle the molecule that corresponds to the spectrum shown.



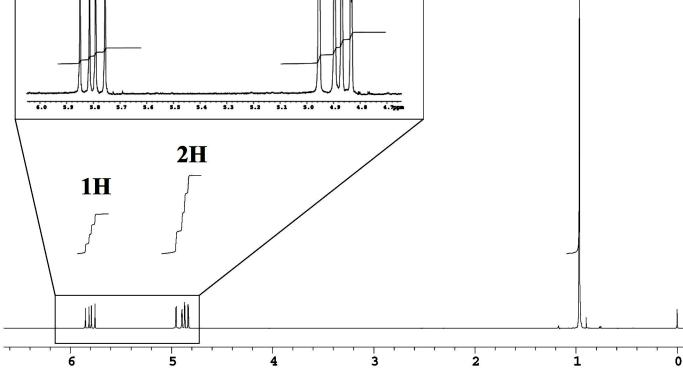




**NOTE:** This box is a magnification of the area below so you can see the signals more clearly!



**9H** 



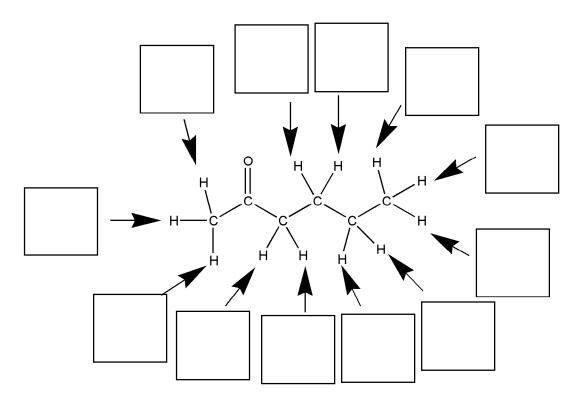
Signature	
-----------	--

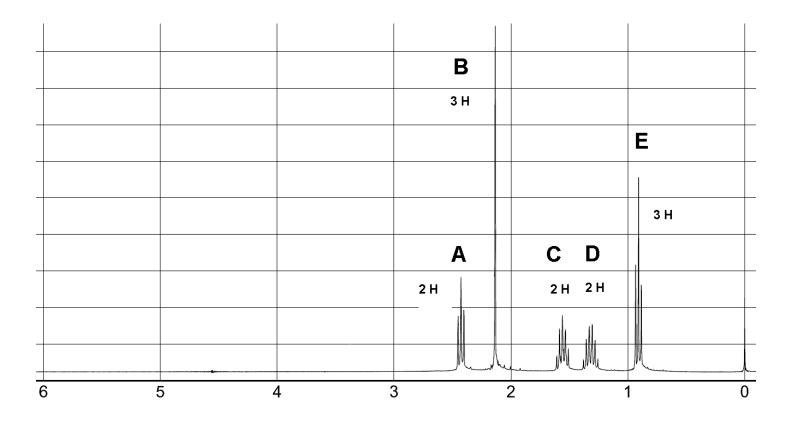
Pg 11

\_(12)

**14.** (12 pts) The following <sup>1</sup>H-NMR spectrum of 2-hexanone has signals labeled with letters. In the boxes provided on the structure, place the letter of the signal that corresponds to the H atoms indicated by the arrows. Because of equivalence, more than one box can get the same letter!

(No D<sub>2</sub>O was added to the sample)





15. (14 pts.) Complete the mechanism for the following reaction of an aldehyde with HCN. Be sure to show arrows to indicate movement of <u>all</u> electrons, write <u>all</u> lone pairs, <u>all</u> formal charges, and <u>all</u> the products for each step. Remember, I said <u>all</u> the products for each step. IF A NEW CHIRAL CENTER IS CREATED IN AN INTERMEDIATE, MARK IT WITH AN ASTERISK. IF A CHIRAL CENTER IS CREATED IN THE PRODUCTS YOU NEED TO DRAW BOTH ENANTIONMERS, AND LABEL THE PRODUCT MIXTURE AS RACEMIC IF RELEVANT. I realize these directions are complex, so please read them again to make sure you know what we want.

HCN Reacting with an Aldehyde

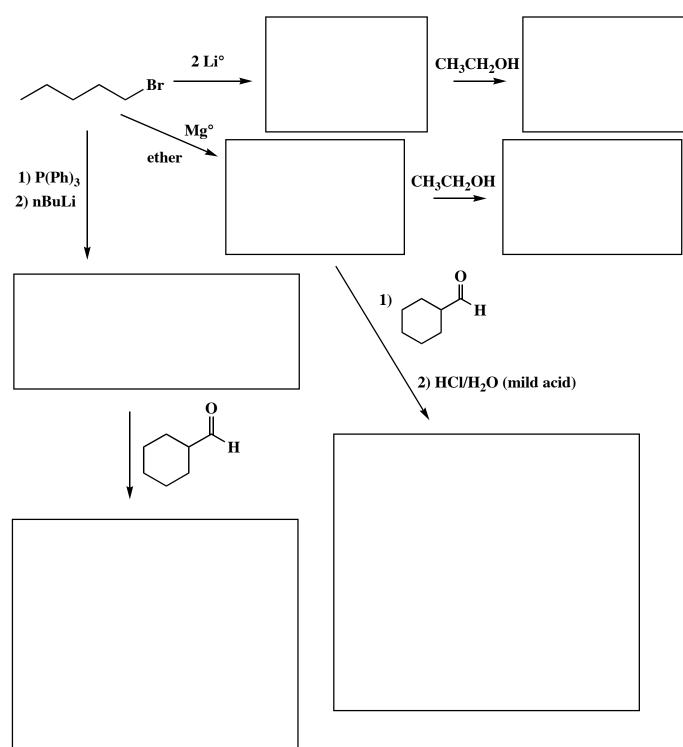
Actual reaction:

Products

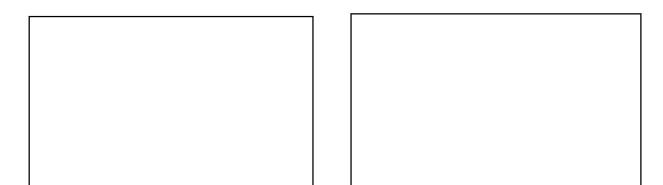
16. (35 pts.) Complete the mechanism for the following acetal formation reaction. Be sure to show arrows to indicate movement of <u>all</u> electrons, write <u>all</u> lone pairs, <u>all</u> formal charges, and <u>all</u> the products for each step. Remember, I said <u>all</u> the products for each step. IF A NEW CHIRAL CENTER IS CREATED IN AN INTERMEDIATE, MARK IT WITH AN ASTERISK. IF A CHIRAL CENTER IS CREATED IN THE PRODUCTS YOU NEED TO DRAW BOTH ENANTIONMERS, AND LABEL THE PRODUCT MIXTURE AS RACEMIC IF RELEVANT. I realize these directions are complex, so please read them again to make sure you know what we want.

Acid Catalyzed Acetal Formation from a Ketone

17. (3 or 5 pts.) Write the predominant product or products that will occur for each transformation. If a new chiral center is created and a racemic mixture is formed, you must draw both enantiomers and write "racemic" under the structure. Use wedges ( ) and dashes ( ) to indicate stereochemistry. To get full credit, you only need to write the major organic product for these. You do not have to worry about the other products.



18. (3 or 5 pts.) Write the predominant product or products that will occur for each transformation. If a new chiral center is created and a racemic mixture is formed, you must draw both enantiomers and write "racemic" under the structure. Use wedges ( ) and dashes ( ) to indicate stereochemistry. To get full credit, you only need to write the major organic product for these. You do not have to worry about the other products.



$$\begin{array}{c|c} O \\ \hline \\ HO \\ \hline \\ HCl \\ (catalytic \\ amount) \end{array} \qquad \begin{array}{c} I) \ CO_2 \\ \hline \\ 2) \ HCl \ / \\ H_2O \\ (strong \\ acid \\ and \\ excess \\ H_2O) \end{array}$$

Signature				

Pg 16\_\_\_\_\_(10)

19. These are synthesis questions. You need to show how the starting material can be converted into the product(s) shown. You may use any reactions we have learned. Show all the reagents you need. Show each molecule synthesized along the way and be sure to pay attention to the regiochemistry and put an asterisk (\*) next to any new chiral center created along the way. You do NOT need to use wedges and dashes and you do NOT need to draw both enantiomers, just mark any new chiral center with an asterisk (\*) AND write "racemic" if appropriate.

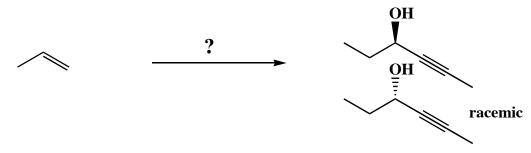
(10 pts) All of the carbon atoms of the products must come from the starting materials for this one!

<b>Signature</b>			

Pg 17\_\_\_\_\_(13)

**19. (cont.)** These are synthesis questions. You need to show how the starting material can be converted into the product(s) shown. You may use any reactions we have learned. Show all the reagents you need. Show each molecule synthesized along the way and be sure to pay attention to the regiochemistry and put an asterisk (\*) next to any new chiral center created along the way. You do NOT need to use wedges and dashes and you do NOT need to draw both enantiomers, just mark any new chiral center with an asterisk (\*) AND write "**racemic**" if appropriate.

(13 pts) All of the carbon atoms of the products must come from the starting materials for this one!



Signature
-----------

Pg 18\_\_\_\_\_(13)

**19.** (cont.) These are synthesis questions. You need to show how the starting material can be converted into the product(s) shown. You may use any reactions we have learned. Show all the reagents you need. Show each molecule synthesized along the way and be sure to pay attention to the regiochemistry and put an asterisk (\*) next to any new chiral center created along the way. You do NOT need to use wedges and dashes and you do NOT need to draw both enantiomers, just mark any new chiral center with an asterisk (\*) AND write "racemic" if appropriate.

(10 pts) All of the carbon atoms of the products must come from the starting materials for this one!

## SAVE THESE UNTIL THE END

**20.** (**5 pts. each**) A fundamental principle of organic chemistry is that functional groups react the same in complex molecules as they do in simple ones. The following two Wittig reactions use more complicated molecules than you are accustomed to seeing, but they react the same. Draw the product of each reaction, making sure to indicate stereochemistry with wedges and dashes.

21. (9 pts.) Here is an APPLY WHAT YOU KNOW QUESTION. You have not seen this before, but with a little information and your background in organic chemistry, we think you can figure it out. Here is what you do not know. Sulfonium ylides can be formed that are analogous to phosphonium ylides (Wittig reagents). However, the sulfonium ylides react with carbonyl compounds to give epoxides. For the following reaction (which you have never seen) predict the intermediate and flow of electrons that leads to the epoxide product. Draw arrows that indicate the flow of all electrons, and be sure to show all lone pairs and formal charges in your intermediate and put an asterisk "\*" next to any new chiral center created.