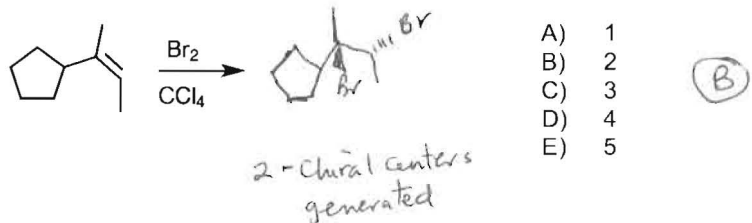


Name: Answer Key

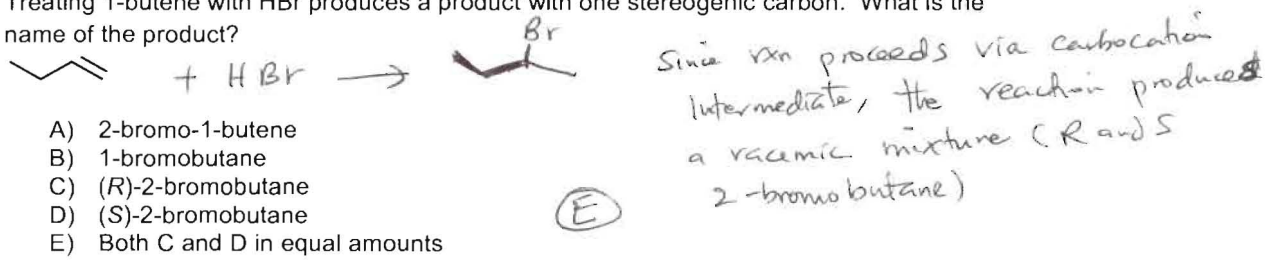
Section _____

CHEM 226: Elementary Organic Chemistry
Exam 2 spring 2013 **March 20, 2013**

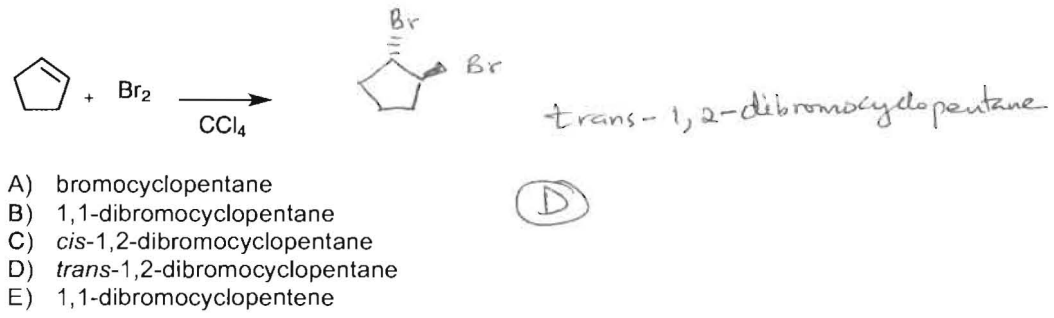
1. How many stereogenic carbons are produced in the product of the following sequence of reactions?



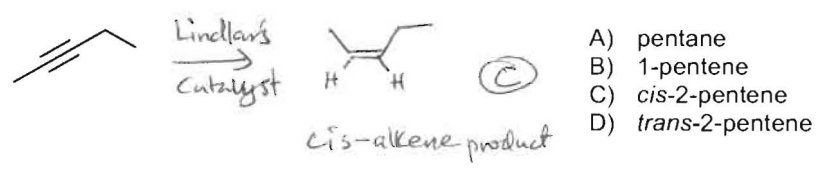
2. Treating 1-butene with HBr produces a product with one stereogenic carbon. What is the name of the product?



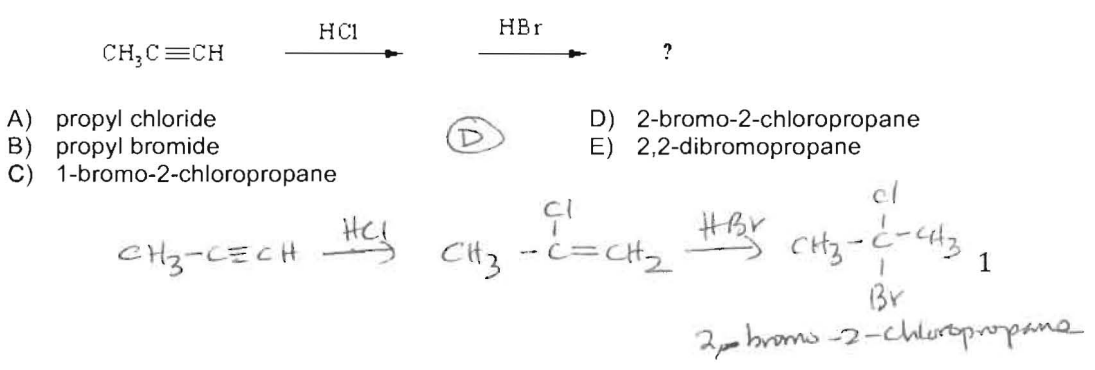
3. What is the name of the product formed from the following reaction?



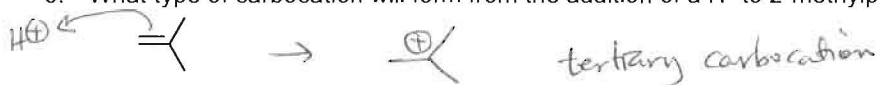
4. Addition of H₂ to 2-pentyne in the presence of the Lindlar's catalyst will produce:



5. What is the final product of adding 1 mole of each reactant in the following sequence?



6. What type of carbocation will form from the addition of a H^+ to 2-methylpropene?



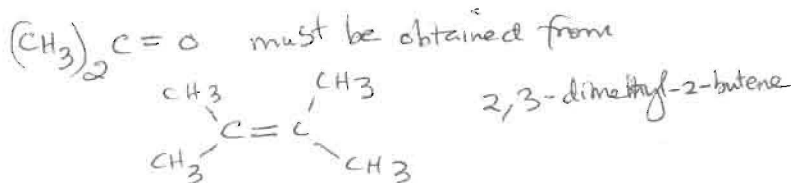
- A) H_3C^+
- B) 1°
- C) 2°
- D) 3°
- E) Allyl

(D)

7. Upon ozonolysis which alkene will give only acetone, $(CH_3)_2C=O$?

- A) 2,3-dimethyl-2-butene
- B) 2,2-dimethyl-2-butene
- C) 3-hexene
- D) 2-methyl-2-pentene
- E) 2-methyl-3-hexene

(A)



8. What is the percent s character in an sp^2 hybrid orbital?

- A) 25%
- B) 33%
- C) 50%
- D) 67%
- E) 75%

s character in $sp^2 = \frac{1}{3} \times 100 = 33\%$

(B)

9. The triple bond in ethyne is made up of

- A) two pi bonds and a sigma bond, each formed by a lateral overlap of two p orbitals.
- B) a sigma bond formed by overlap of two s orbitals and two pi bonds, each formed by lateral overlap of two p orbitals.
- (D) C) a sigma bond formed by end-on overlap of two sp^2 orbitals and a pi bond formed by lateral overlap of two p orbitals.
- D) two pi bonds, each formed by lateral overlap of two p orbitals, and a sigma bond formed by end-on overlap of two sp orbitals. ✓
- E) two pi bonds, each formed by end-on overlap of two p orbitals, and a sigma bond formed by lateral overlap of two sp orbitals.

10. Bending vibrations in the infrared region occur at:

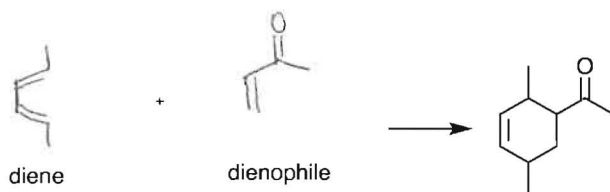
- A) 3000 cm^{-1}
- B) 2200 cm^{-1}
- C) 1700 cm^{-1}
- D) below 1400 cm^{-1}
- E) over 3000 cm^{-1}

often shows in the fingerprint region below 1400 cm^{-1}

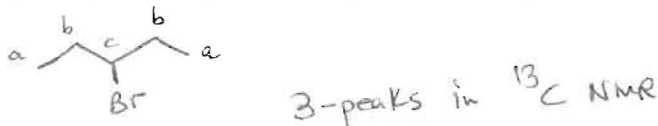
(D)

Short Answer Questions

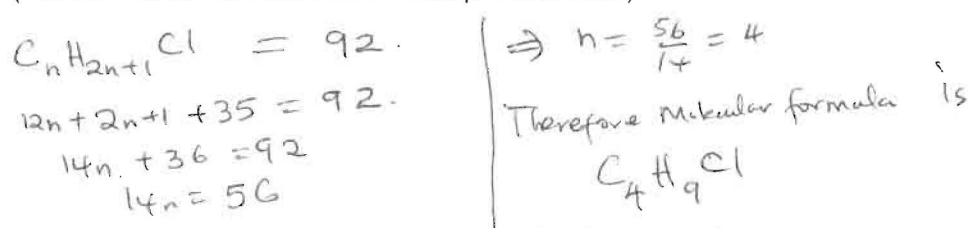
11. The Diels-Alder reaction is very important in the synthesis of six-membered rings. Draw the reagents that can be used to synthesize the product shown by this method?



12. How many peaks would you expect in the proton decoupled ^{13}C NMR spectrum of 3-bromopentane?

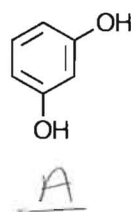
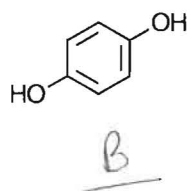


13. A monochloroalkane shows two parent ion peaks m/z at 92 and 94. What is the molecular formula? (^{35}Cl and ^{37}Cl are the most common isotopes of chlorine)



14. A student lost the labels of two compounds and was required to run experiments to distinguish and identify them. She took an IR spectrum of both compounds A and B and both showed a broad band in the 3200 to 3500 cm^{-1} region of their IR spectrum? She then took both ^1H NMR and ^{13}C NMR spectra, and compound A had four peaks, while compound B had two peaks in both NMRs. Identify compounds A and B.

Both had -OH groups.



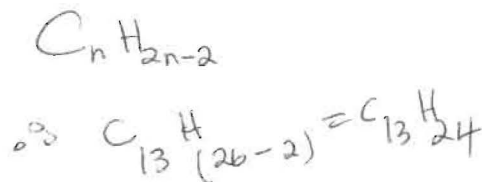
A - 4 peaks ^1H NMR
 ^{13}C NMR

B - 2 peaks ^1H NMR
 ^{13}C NMR

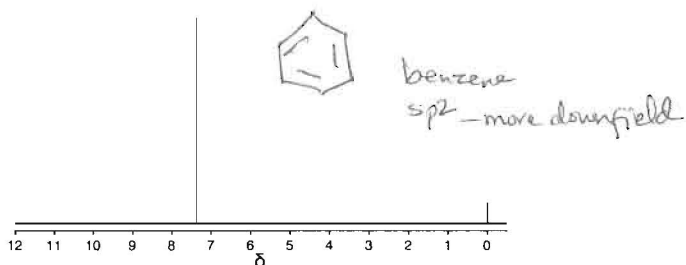
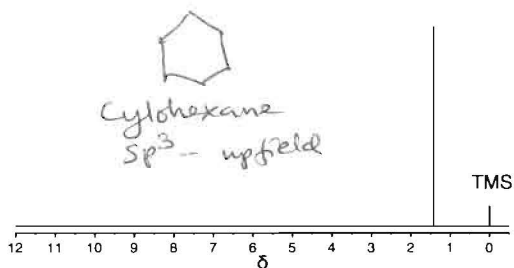
15. What is the molecular geometry of an alkynes like acetylene and a hydrogen cyanide
 $\text{HC}\equiv\text{N}$ $\text{HC}\equiv\text{CH}$



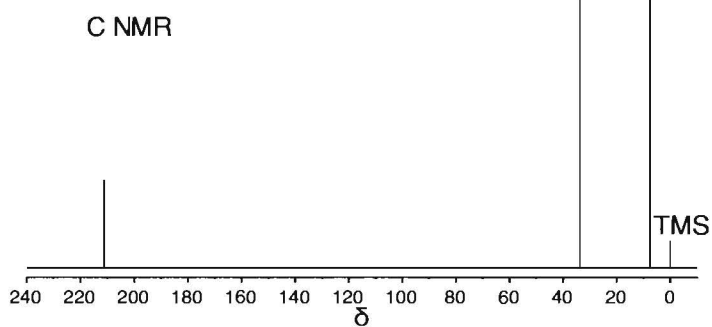
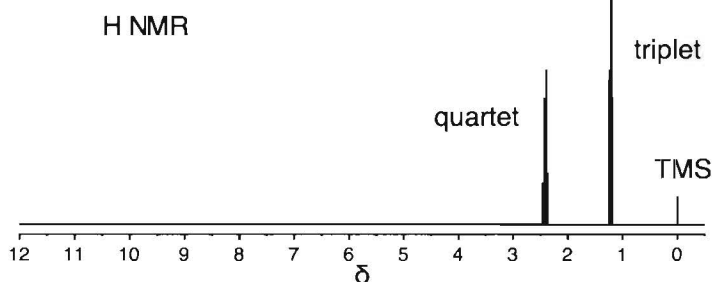
16. How many hydrogens are there in an alkyne with 13 carbons? 24



17. The ^1H NMR spectrum of benzene (C_6H_6) and cyclohexane (C_6H_{12}) are shown below. Match the spectra with the correct molecule.



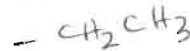
18. A compound, $\text{C}_5\text{H}_{10}\text{O}$, has an intense IR band at 1725 cm^{-1} . Its ^1H NMR and ^{13}C NMR are shown below. Deduce the structure of the compound.



•) strong IR band at 1725 cm^{-1} suggests a carbonyl group

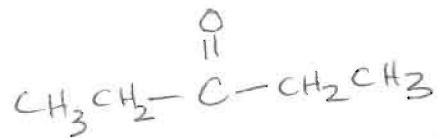


•) triplet, quartet pattern in ^1H NMR suggests an ethyl group



• Three peaks in ^{13}C NMR suggests three types of carbons

from the molecular formula and the structure of the compound is



For questions 19-25, fill in the missing reagent(s) and the major product(s) for the following reactions

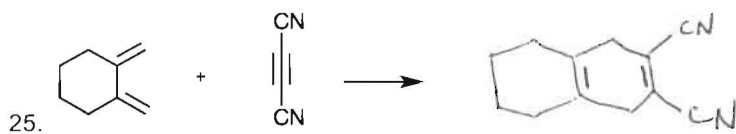
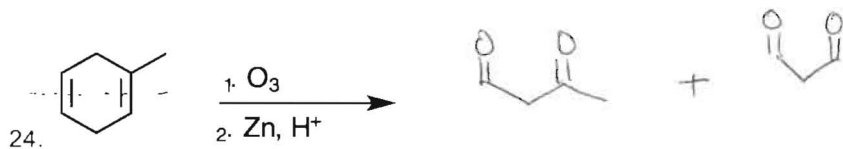
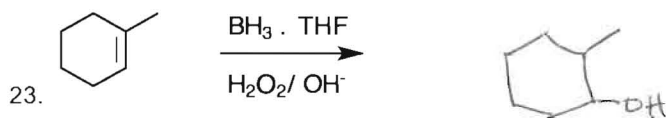
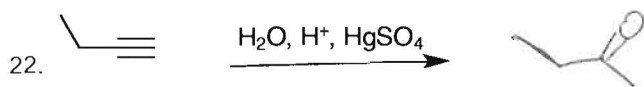
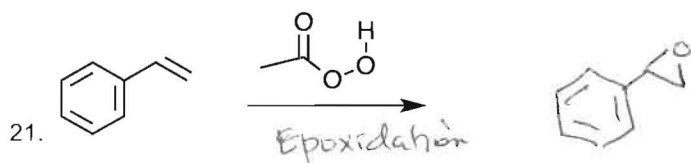
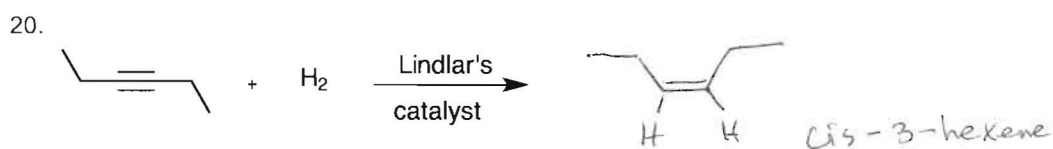


Table 12.2 Typical ^1H Chemical Shifts (Relative to Tetramethylsilane)

Type of ^1H	δ (ppm)	Type of ^1H	δ (ppm)
$\text{C}-\text{CH}_3$	0.85-0.95	$-\text{CH}_2-\text{F}$	4.3-4.4
$\text{C}-\text{CH}_2-\text{C}$	1.20-1.35	$-\text{CH}_2-\text{Br}$	3.4-3.6
		$-\text{CH}_2-\text{I}$	3.1-3.3
$\begin{array}{c} \text{C} \\ \\ \text{C}-\text{CH}-\text{C} \end{array}$	1.40-1.65	$\text{CH}_2=\text{C}$	4.6-5.0
$\text{CH}_3-\text{C}=\text{C}$	1.6-1.9	$-\text{CH}=\text{C}$	5.2-5.7
CH_3-Ar	2.2-2.5	$\text{Ar}-\text{H}$	6.6-8.0
$\begin{array}{c} \text{CH}_3-\text{C}=\text{O} \\ \end{array}$	2.1-2.6	$-\text{C}\equiv\text{C}-\text{H}$	2.4-2.7
$\text{CH}_3-\text{N} \begin{array}{l} \diagup \\ \diagdown \end{array}$	2.1-3.0	$\begin{array}{c} \text{O} \\ \\ -\text{C}-\text{H} \end{array}$	9.5-9.7
$\text{CH}_3-\text{O}-$	3.5-3.8	$\begin{array}{c} \text{O} \\ \\ -\text{C}-\text{OH} \end{array}$	10-13
$-\text{CH}_2-\text{Cl}$	3.6-3.8	$\text{R}-\text{OH}$	0.5-5.5
$-\text{CHCl}_2$	5.8-5.9	$\text{Ar}-\text{OH}$	4-8

Table 12.4 Infrared Stretching Frequencies of Some Typical Bonds

Bond type	Group	Class of compound	Frequency range (cm^{-1})
single bonds to hydrogen	$\text{C}-\text{H}$	alkanes	2850-3000
	$=\text{C}-\text{H}$	alkenes and aromatic compounds	3030-3140
	$\equiv\text{C}-\text{H}$	alkynes	3300
	$\text{O}-\text{H}$	alcohols and phenols	3500-3700 (free) 3200-3500 (hydrogen-bonded)
	$\text{O}-\text{H}$	carboxylic acids	2500-3000
	$\text{N}-\text{H}$	amines	3200-3600
double bonds	$\text{S}-\text{H}$	thiols	2550-2600
	$\text{C}=\text{C}$	alkenes	1600-1680
	$\text{C}=\text{N}$	imines, oximes	1500-1650
	$\text{C}=\text{O}$	aldehydes, ketones, esters, acids	1650-1780
triple bonds	$\text{C}\equiv\text{C}$	alkynes	2100-2260
	$\text{C}\equiv\text{N}$	nitriles	2200-2400