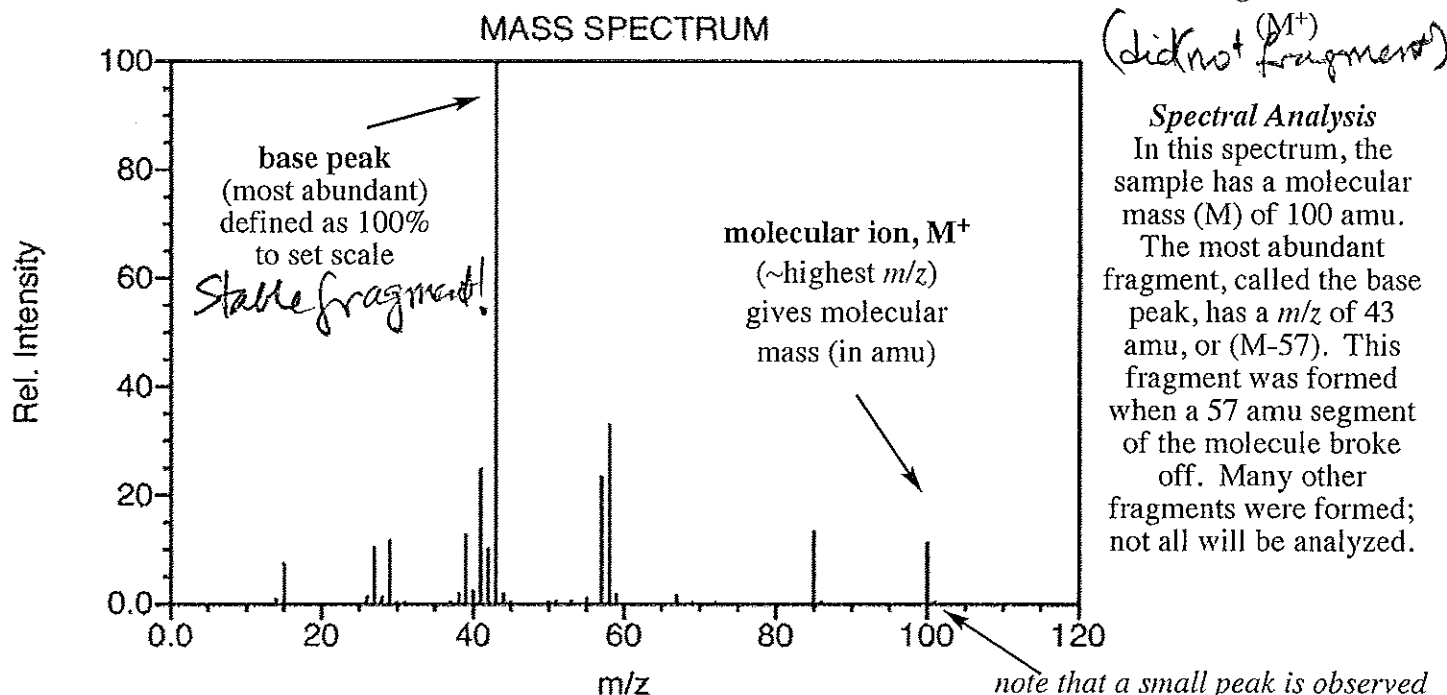
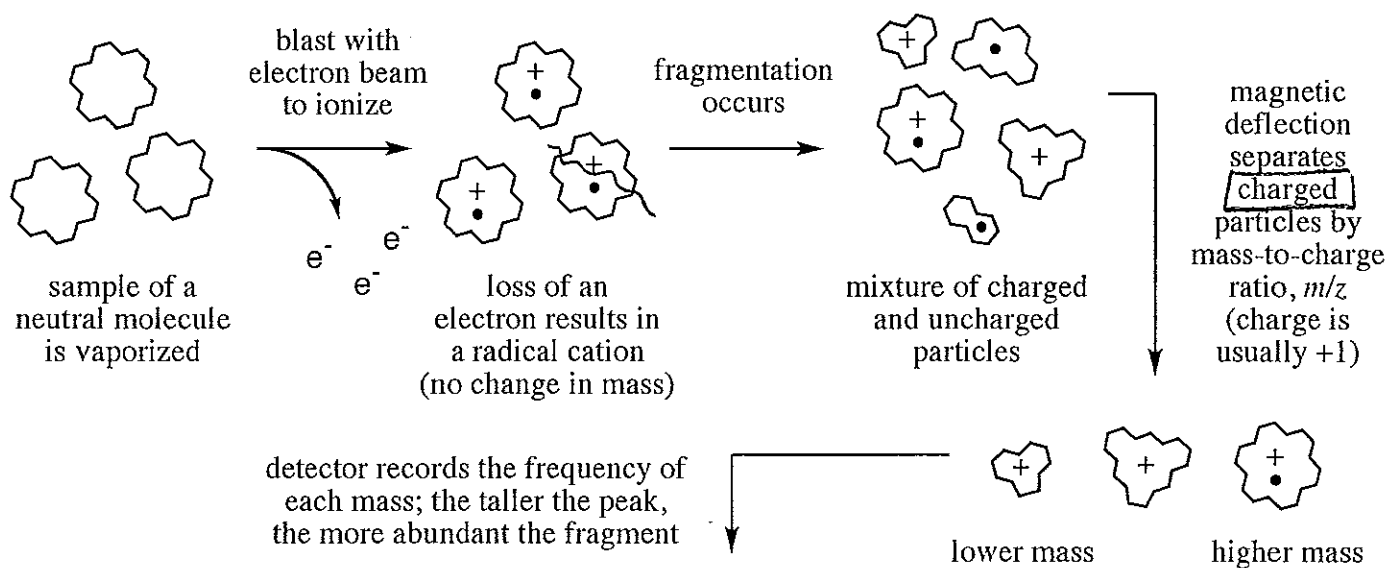


CHM 319L Organic Laboratory, Dr. Laurie S. Starkey

Introduction to Mass Spectrometry

Mass spectrometry is used to determine a sample's molecular mass and molecular formula. Some structural information can also be determined by mass spec. This technique is especially valuable when used in conjunction with gas chromatography (called GC-MS); the GC separates the components of a mixture and then the mass spec analyzes each component. To obtain a mass spectrum, a sample is vaporized and then bombarded with a high energy beam of electrons (this technique is called electron-impact ionization, or EI). This ionizes the sample by knocking an electron out of the structure. A nonbonded or bonded electron is removed, resulting in radical cation species. Although it is now missing an electron, the intact molecule still has the same molecular mass and is known as the **molecular ion (M^+)**, or the parent ion. In addition, this high-energy environment causes the molecular ion to fragment and the various pieces can be analyzed to learn something about the original structure. All **charged species** are separated by mass as they are passed through a magnetic field and a detector records how many pieces of each mass have been formed. Note that because this technique does not involve the absorption or emission of energy, it is not called spectroscopy.



spectra have been reproduced from the NIST Chemistry WebBook <http://webbook.nist.gov/chemistry>

What is the mass of a single molecule?

CH_4 Mass = C + 4(H) = 12 + 4(1) = 16 amu (M^+)

but ~1% of carbon atoms exist as ^{13}C isotope!

$^{13}\text{CH}_4$ Mass = ^{13}C + 4(H) = 13 + 4(1) = 17 amu ($M+1$) peak
 ~1% of M^+ peak

What ratio is expected for the molecular ion peaks of C_2H_6 ?

C_2H_6 or $^{13}\text{CH}_3-\text{CH}_3$ or $\text{CH}_3-^{13}\text{CH}_3$ $2\text{C} \rightarrow (M+1)$ is 2% of M

the number of carbon atoms affects the relative height of the $M+1$ peak

Other isotopes of high abundance

Cl (35.45 amu) atoms are 76% ^{35}Cl and 24% ^{37}Cl ~ 3:1 $M : M+2$

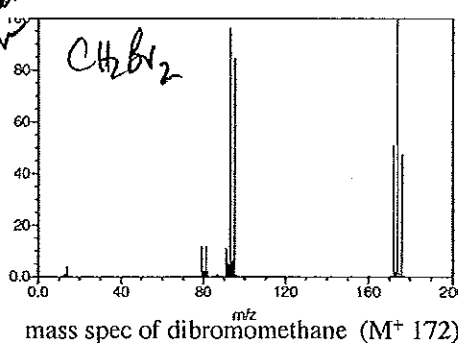
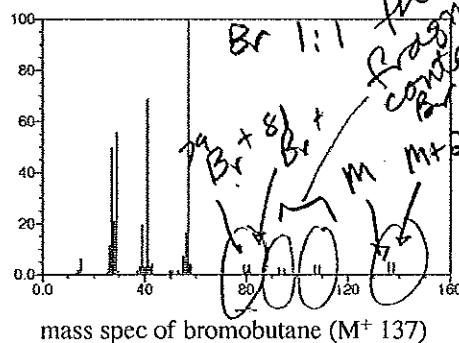
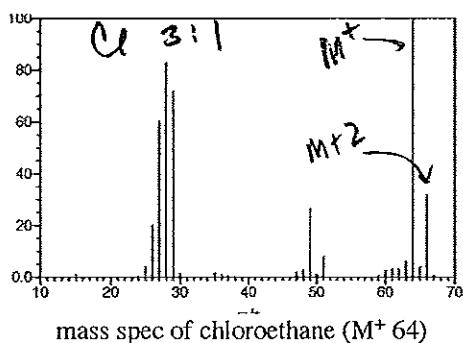
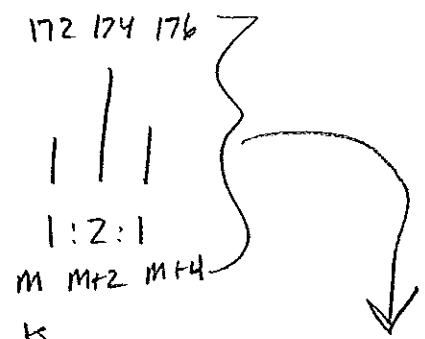
Br (79.90 amu) atoms are 51% ^{79}Br and 49% ^{81}Br ~ 1:1 $M : M+2$

fragments containing Cl or Br have significant $M+2$ peaks

What ratios are expected for the molecular ion peaks of CH_2Br_2 ?

possible masses?	CH_2	Br	Br	total amu
↓	14	79	79	172
four combinations	14	81	79	174
	14	79	81	174
	14	81	81	176

twice as likely as other combos



Determining molecular formula from high-resolution mass spectrometry (high-res mass spec)

exact masses of various elements:

^1H	1.007825 amu
^{12}C	12.000000 amu
^{14}N	14.003050 amu
^{16}O	15.994914 amu

these molecules all have a molecular mass of 98 g/mol

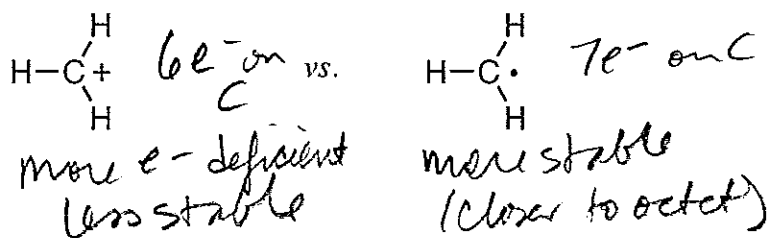
$\text{C}_4\text{H}_6\text{N}_2\text{O}$	98.0480 amu
$\text{C}_5\text{H}_6\text{O}_2$	98.0368 amu
$\text{C}_5\text{H}_{10}\text{N}_2$	98.0845 amu
$\text{C}_6\text{H}_{10}\text{O}$	98.0732 amu
C_7H_{14}	98.1096 amu

the molecular formula can be determined from precise molecular mass

Mass Spec Features of Various Functional Groups

Fragmentation of the molecular ion

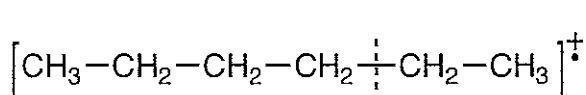
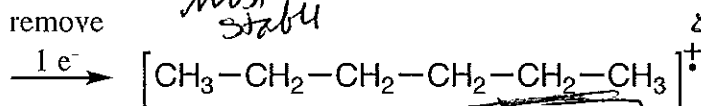
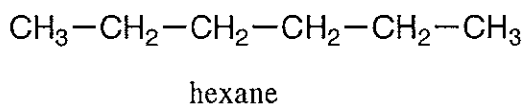
Which is more stable, a carbocation C⁺ or a radical R• ?



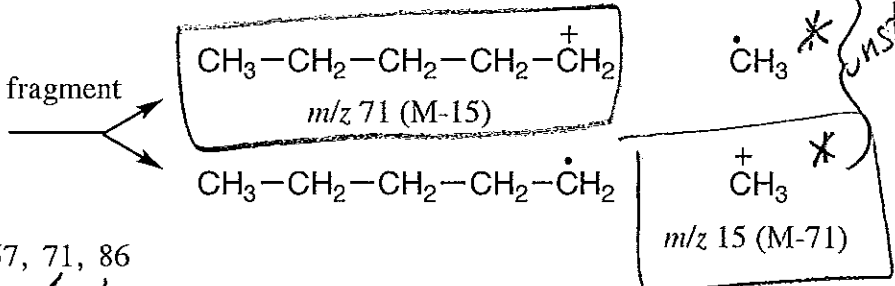
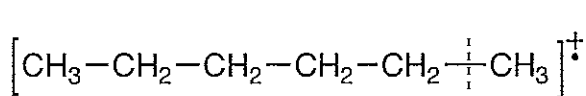
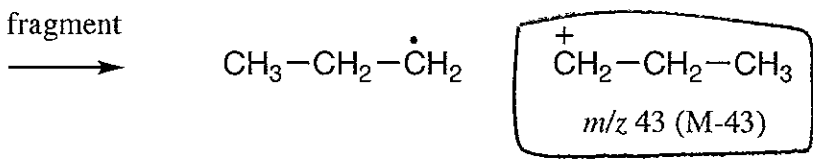
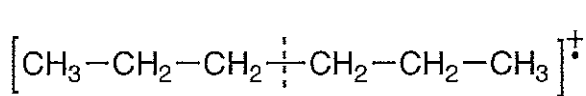
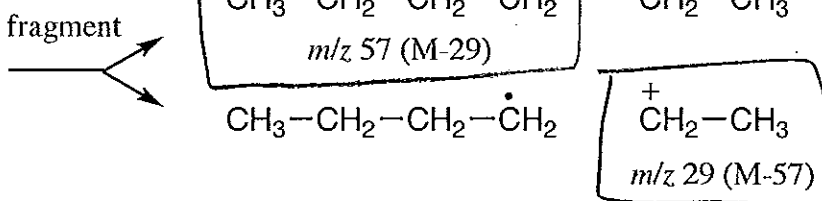
fragmentation is more likely if it gives more stable carbocations and radicals

3° > 2° > 1° > methyl
 most stable vs. least stable

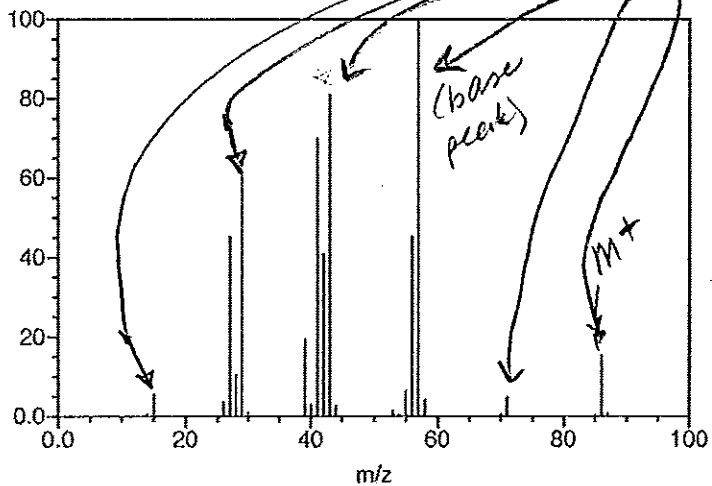
Alkanes



break a bond to give two fragments: one cation and one radical

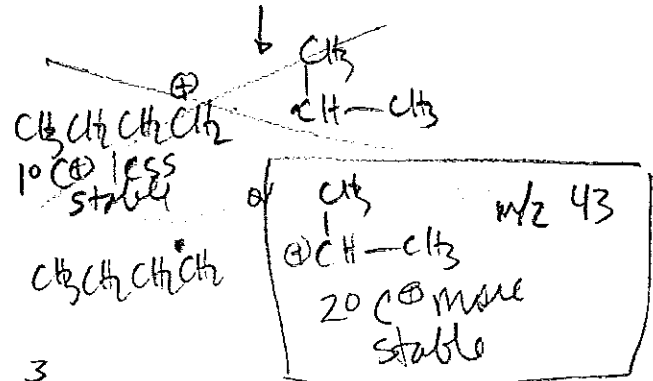
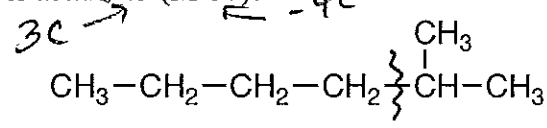


note: hexane has peaks at m/z 15, 29, 43, 57, 71, 86



Branched Alkanes

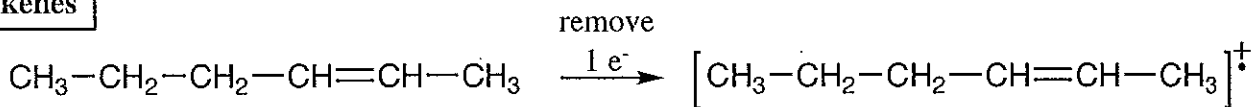
Explain why the base peak of 2-methylhexane is at m/z 43 (M-57).



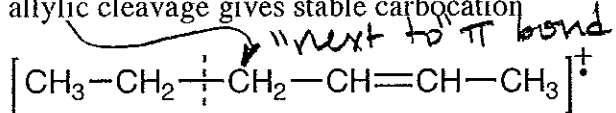
Why are the peaks at m/z 15 and 71 so small?

* This fragmentation is less likely since it results in less stable methyl radical + carbocation

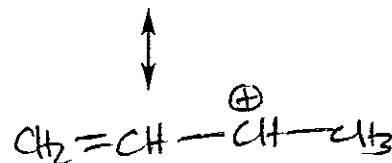
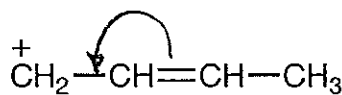
Alkenes



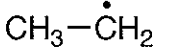
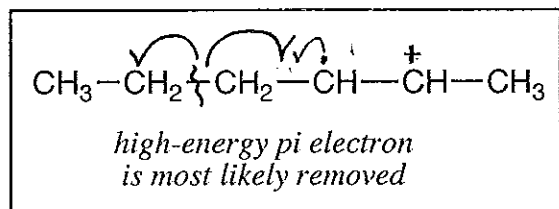
allylic cleavage gives stable carbocation



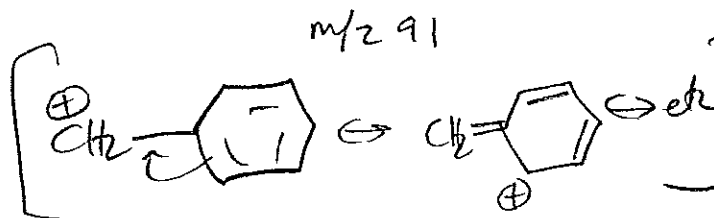
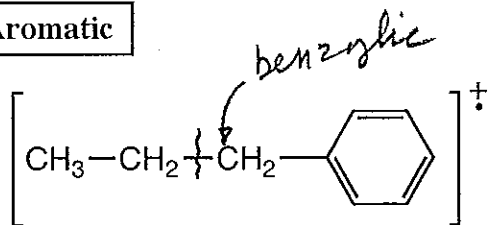
fragment



allylic carbocation is resonance-stabilized

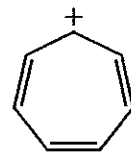


Aromatic



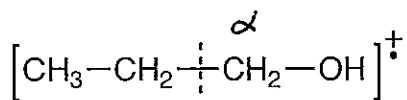
benzylic \oplus resonance

rearranges

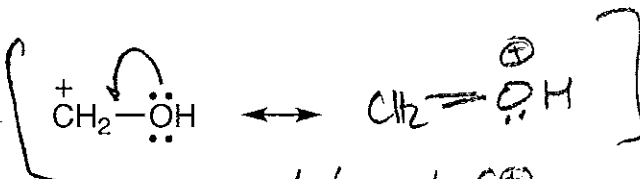


tropylium cation
m/z 91

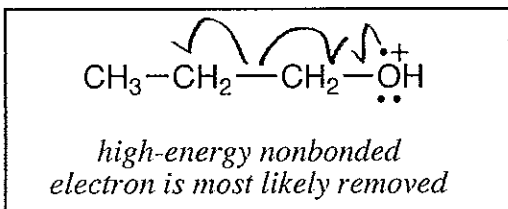
Alcohols



α cleavage

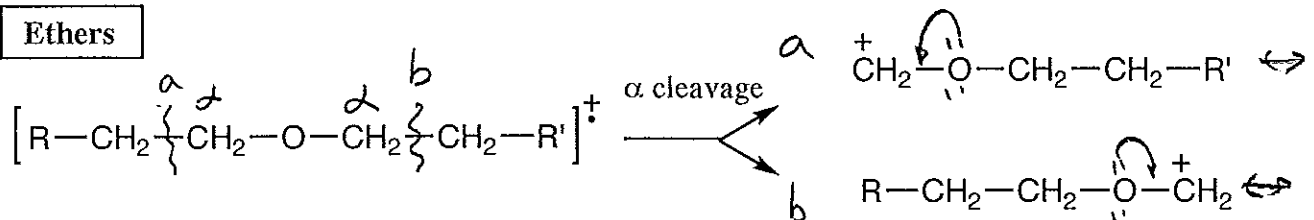


res stabilized \oplus

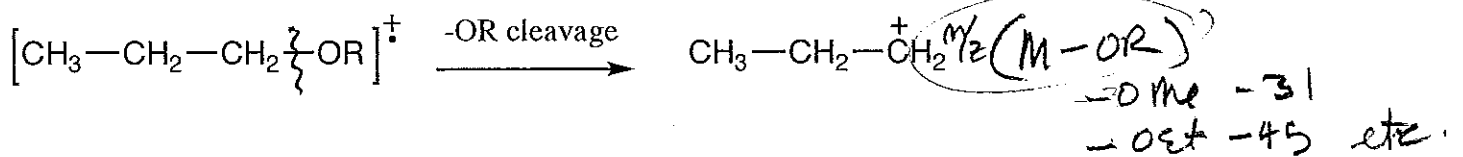


*also, loss of H₂O (M-18) may be observed
(multistep mechanism first involves an intramolecular proton transfer)

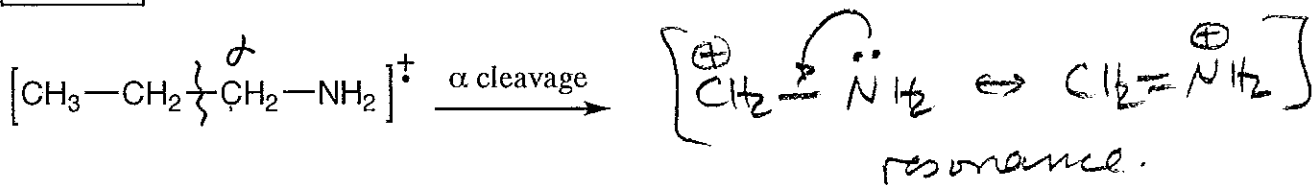
Ethers



also, loss of RO• may be observed



Amines

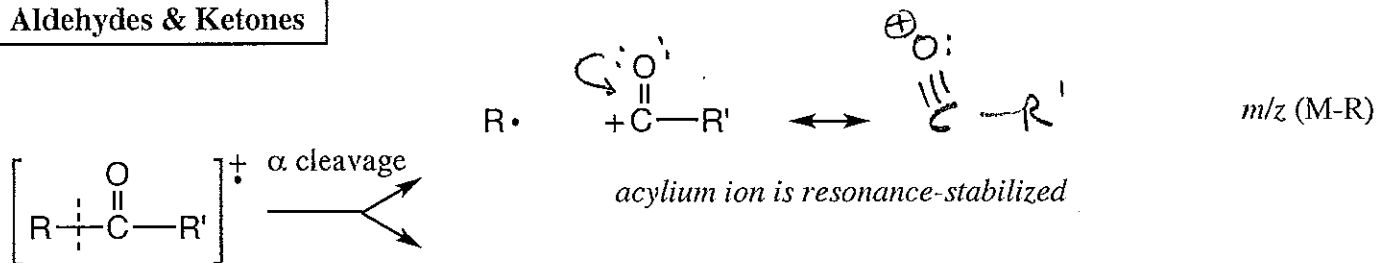


* **Nitrogen Rule:** M^+ is odd if there are an odd number of nitrogens!

odd molecular mass indicates one nitrogen (or 3 or 5 or 7 nitrogens...)

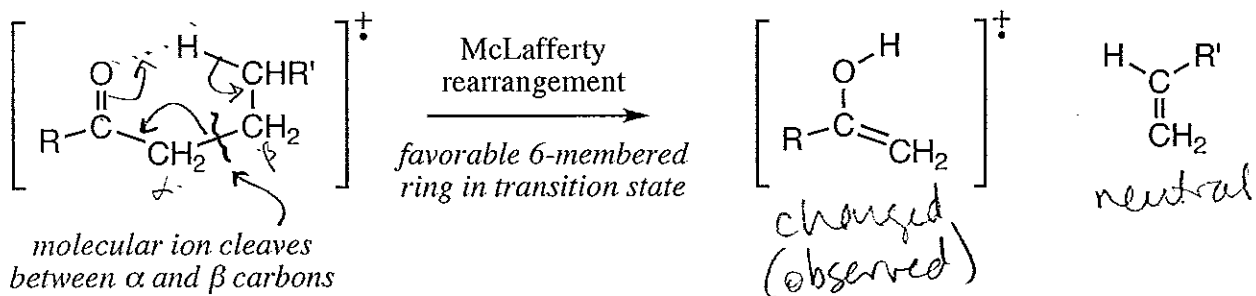
even molecular mass indicates zero nitrogens (or 2 or 4 or 6 nitrogens...)

Aldehydes & Ketones



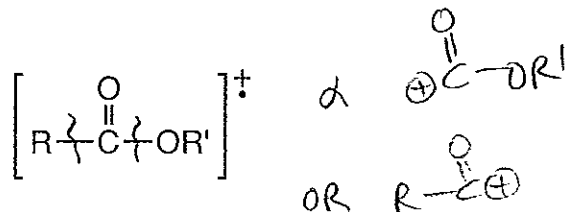
loss of larger group more likely
(more stable R•)

McLafferty rearrangement is a common fragmentation for carbonyl-containing molecules



Esters

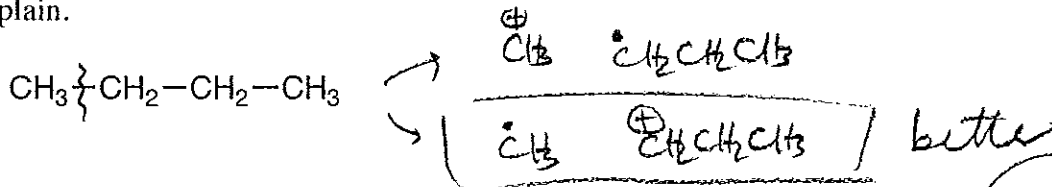
esters can undergo α cleavage, McLafferty rearrangement, or loss of RO•



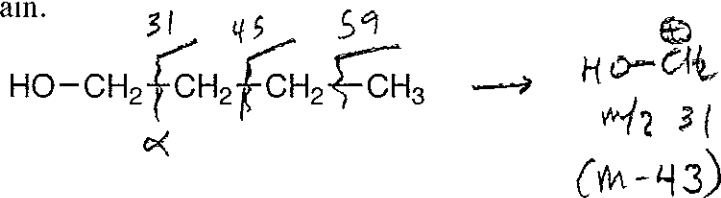
work 25-30 minutes on their own (or in groups) methyl propyl

Mass Spectrometry Discussion Questions

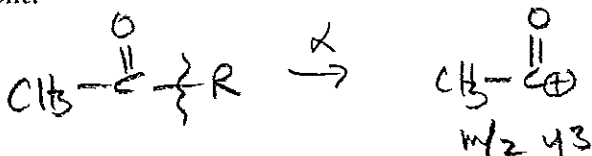
For the given molecule ($M=77$), do you expect the more abundant peak to be m/z 15 or m/z 43? Explain.



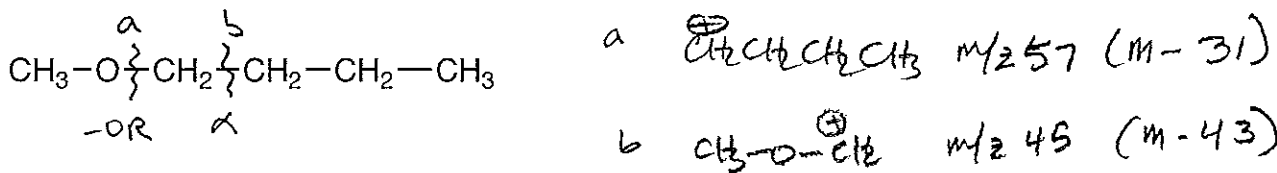
For the given molecule ($M=74$), which peak do you expect to be most abundant: m/z 31, m/z 45 or m/z 59? Explain.



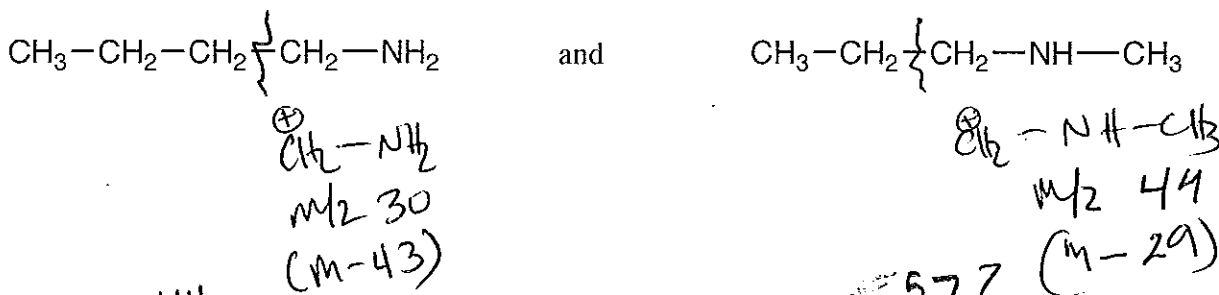
Explain why the mass spectra of methyl ketones typically have a peak at m/z 43. Provide the structure of this fragment.



In the mass spectrum of the given molecule ($M=88$), account for the peaks at m/z 45 and m/z 57.



How could you use mass spectrometry to distinguish between the following two compounds ($M=73$)? Provide structures (and m/z values) for the significant fragments expected.



What would be the m/z ratio for the fragment resulting from a McLafferty Rearrangement for the following molecule ($M=100$)? What fragment accounts for its base peak at m/z 42?

