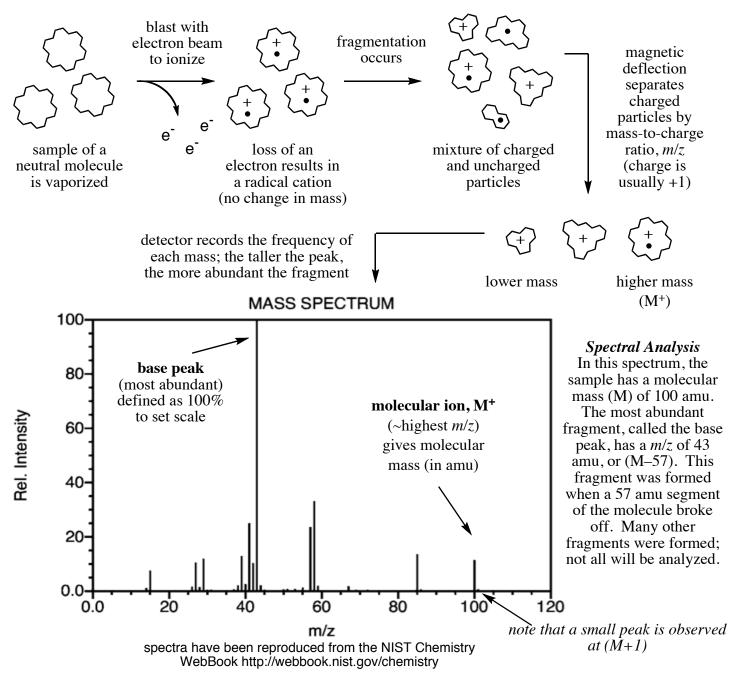
CHM 319L Organic Laboratory, Dr. Laurie S. Starkey Introduction to Mass Spectrometry

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 conjuction 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.



What is the mass of a single molecule?

$$CH_4$$
 Mass = $C + 4(H) =$

but $\sim 1\%$ of carbon atoms exist as 13 C isotope!

$$^{13}CH_4$$
 Mass = $^{13}C + 4(H) =$

What ratio is expected for the molecular ion peaks of C₂H₆?

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

Other isotopes of high abundance

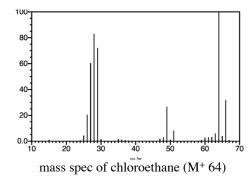
Cl (35.45 amu) atoms are 76% 35 Cl and 24% 37 Cl

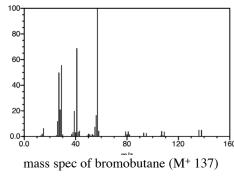
Br (79.90 amu) atoms are 51% ^{79}Br and 49% ^{81}Br

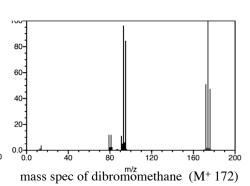
fragments containing Cl or Br have significant M+2 peaks

What ratios are expected for the molecular ion peaks of CH₂Br₂?

possible masses? CH₂ Br Br total amu







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

exact masses of various elements:

¹H 1.007825 amu
 ¹²C 12.000000 amu
 ¹⁴N 14.003050 amu

¹⁴N 14.003050 amu ¹⁶O 15.994914 amu these molecules all have a molecular mass of 98 g/mol $\begin{cases} C_4H_6N_2O & 98.0480 \text{ amu} \\ C_5H_6O_2 & 98.0368 \text{ amu} \\ C_5H_{10}N_2 & 98.0845 \text{ amu} \\ C_6H_{10}O & 98.0732 \text{ amu} \\ C_7H_{14} & 98.1096 \text{ amu} \end{cases}$

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

Alkanes

remove $CH_3 - CH_2 - CH_2 - CH_2 - CH_3 \xrightarrow{1 e^-} [CH_3 - CH_2 - CH_2 - CH_2 - CH_2 - CH_3]^{\frac{1}{2}}$

a radical cation (M^+) m/z 86

$$\begin{bmatrix} \mathsf{CH_3} - \mathsf{CH_2} - \mathsf{CH_2} - \mathsf{CH_2} + \mathsf{CH_2} - \mathsf{CH_3} \end{bmatrix}^{\frac{1}{4}} \xrightarrow{\text{fragment}}$$

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

$$\begin{bmatrix} \mathsf{CH_3} - \mathsf{CH_2} - \mathsf{CH_2} + \mathsf{CH_2} - \mathsf{CH_2} - \mathsf{CH_3} \end{bmatrix}^{\frac{1}{\mathsf{CH_2}}} \xrightarrow{\text{fragment}} \\ \mathsf{CH_3} - \mathsf{CH_2} - \dot{\mathsf{CH_2}} + \dot{\mathsf{CH_2}} - \mathsf{CH_2} - \mathsf{CH_2} + \mathsf{CH_2} - \mathsf{CH_3} \end{bmatrix}$$

$$CH_3-CH_2-CH_2-CH_2$$
 $m/z 57 (M-29)$

$$CH_3-CH_2-\dot{C}H_2-\dot{C}H_2$$
 CH_2-CH_3 m/z 29 (M-57)

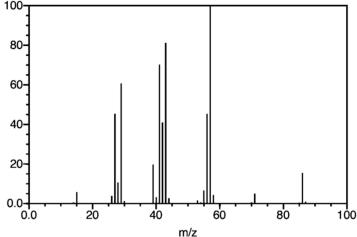
$$CH_3 - CH_2 - \dot{C}H_2$$
 $CH_2 - CH_2 - CH_3$
 $m/z 43 (M-43)$

$$\left[\text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{CH}_2 - \text{CH}_2 - \text{CH}_3 \right]^{\frac{1}{2}} \xrightarrow{\text{fragment}}$$

 $CH_3-CH_2-CH_2-CH_2-CH_2$ СH₃

$${\rm CH_3-CH_2-CH_2-\dot{C}H_2}$$
 ${\rm CH_3 \atop m/z\ 15 \atop (M-71)}$

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



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

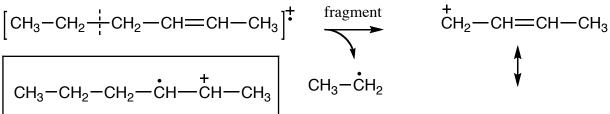
Branched Alkanes

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

$$\begin{array}{c} \operatorname{CH_3} \\ \operatorname{CH_3-CH_2-CH_2-CH_2-CH-CH_3} \end{array}$$

$$CH_3 - CH_2 - CH_2 - CH = CH - CH_3$$
 $\xrightarrow{1 e^-}$ $\left[CH_3 - CH_2 - CH_2 - CH = CH - CH_3 \right]^{\frac{1}{2}}$

allylic cleavage gives stable carbocation



high-energy pi electron is most likely removed

allylic carbocation is resonance-stabilized

Aromatic

$$\begin{bmatrix} CH_3 - CH_2 - CH_2$$



tropylium cation m/z 91

Alcohols

$$\begin{bmatrix} \mathsf{CH_3} - \mathsf{CH_2} + \mathsf{CH_2} - \mathsf{OH} \end{bmatrix}^{\overset{\bullet}{+}} \xrightarrow{\alpha \text{ cleavage}} \quad \overset{\bullet}{\mathsf{CH_2}} - \overset{\bullet}{\mathsf{OH}} \quad \overset{\bullet}{\longleftarrow}$$

$$CH_3-CH_2-CH_2-OH$$

high-energy nonbonded electron is most likely removed

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

Ethers
$$\begin{bmatrix} R-CH_2-CH_2-O-CH_2-CH_2-R' \end{bmatrix}^{\frac{1}{2}} \xrightarrow{\alpha \text{ cleavage}} \begin{bmatrix} CH_2-O-CH_2-CH_2-R' \\ R-CH_2-CH_2-O-CH_2\end{bmatrix}^{\frac{1}{2}}$$

also, loss of RO • may be observed

$$\left[\text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{OR} \right]^{\frac{1}{4}} \quad \xrightarrow{\text{-OR cleavage}} \quad \text{CH}_3 - \text{CH}_2 - \overset{\text{+}}{\text{CH}_2} \qquad \textit{m/z (M-OR)}$$

Amines

$$\begin{bmatrix} CH_3-CH_2-CH_2-NH_2 \end{bmatrix}^{\frac{1}{2}}$$
 $\frac{\alpha \text{ cleavage}}{}$

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

$$R \cdot + C - R' \longrightarrow m/z \text{ (M-R)}$$

$$\begin{bmatrix} O \\ R + C - R' \end{bmatrix} \cdot \alpha \text{ cleavage}$$

$$acylium ion is resonance-stabilized$$

loss of larger group more likely (more stable $R \cdot$)

McLafferty rearrangement is a common fragmentation for carbonyl-containing molecules

molecular ion cleaves between α and β carbons

Esters

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

$$\begin{bmatrix} O \\ R - C - OR' \end{bmatrix}^{\frac{1}{2}}$$

Mass Spectrometry Discussion Questions

For the given molecule (M=58), 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), provide structures 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.

$$CH_3-CH_2-CH_2-CH_2-NH_2$$
 and $CH_3-CH_2-CH_2-NH-CH_3$

What would be the m/z ratio for the fragment resulting from a McLafferty Rearrangement for the following molecule (M=114)? What fragment accounts for its base peak at m/z 57?

$$\begin{array}{c} & \text{O} \\ \text{CH}_{3} - \text{CH} - \text{CH}_{2} - \overset{\text{O}}{\text{C}} - \text{CH}_{2} - \text{CH}_{3} \\ \text{CH}_{3} \end{array}$$

Mass spec problems (Wade 7th, 8th or 9th edition) Chapter 12: 8-11, 18, 20.