

Figure 13.5

$^1\text{H-NMR}$ spectrum of methyl acetate

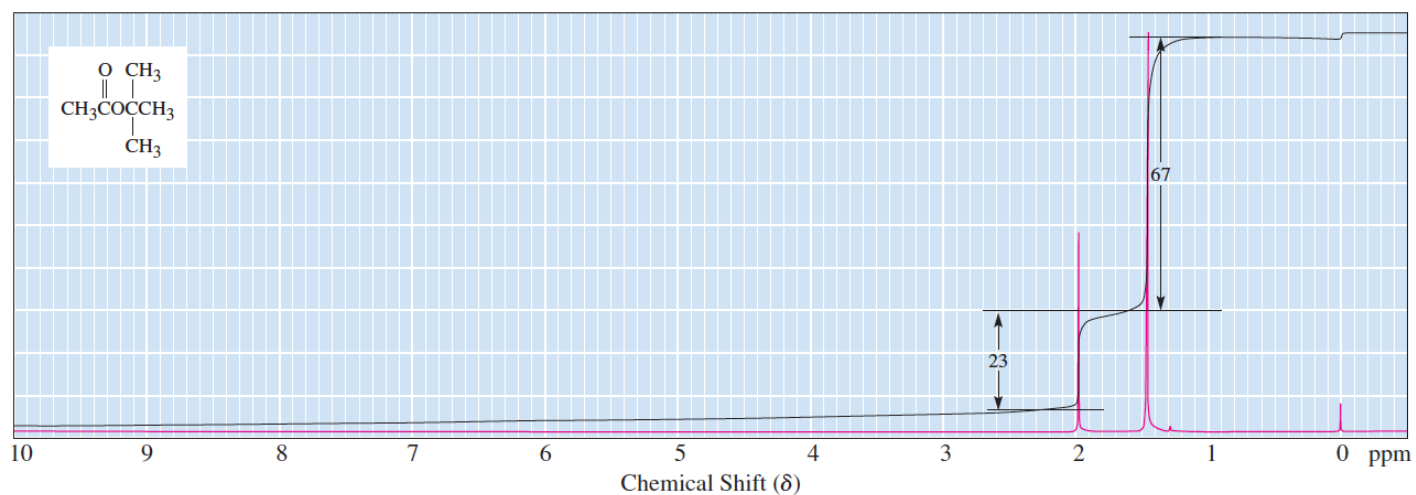


Figure 13.7

$^1\text{H-NMR}$ spectrum of *tert*-butyl acetate showing the integration. The total vertical rise of 90 chart divisions corresponds to 12 hydrogens, 9 in one set and 3 in the other.

J. Adjacent nuclei have magnetic fields associated with their spins. The spins of equivalent adjacent nuclei can be either $+1/2$ or $-1/2$, and at room temperature they are found in about a 50:50 mixture at any given nucleus (very slight excess of lower energy $+1/2$). These can add to give $n+1$ different spin combinations in the proportions predicted by Pascal's triangle. Each different spin combination produces a different magnetic field, which leads to $n+1$ splittings in the peaks of the NMR spectra of the adjacent (no more than three bonds away) nuclei.

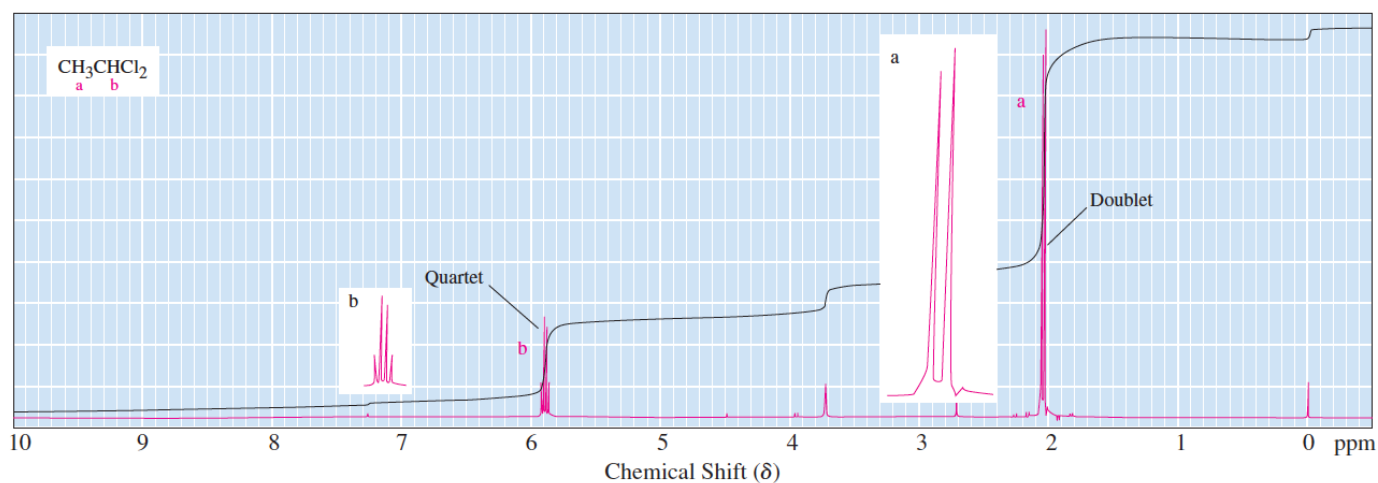


Figure 13.12

$^1\text{H-NMR}$ spectrum of 1,1-dichloroethane.

K. THEORY: When there are two sets of adjacent H atoms, the number of peaks multiply. For example, a CH_2 group with a CH_2 group and a CH_3 group on either side should show $3 \times 4 = 12$ splittings! You can say this group is a "triplet of quartets" (or a "quartet of triplets").

L. WHAT YOU WILL SEE IN REALITY : For alkyl groups 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.

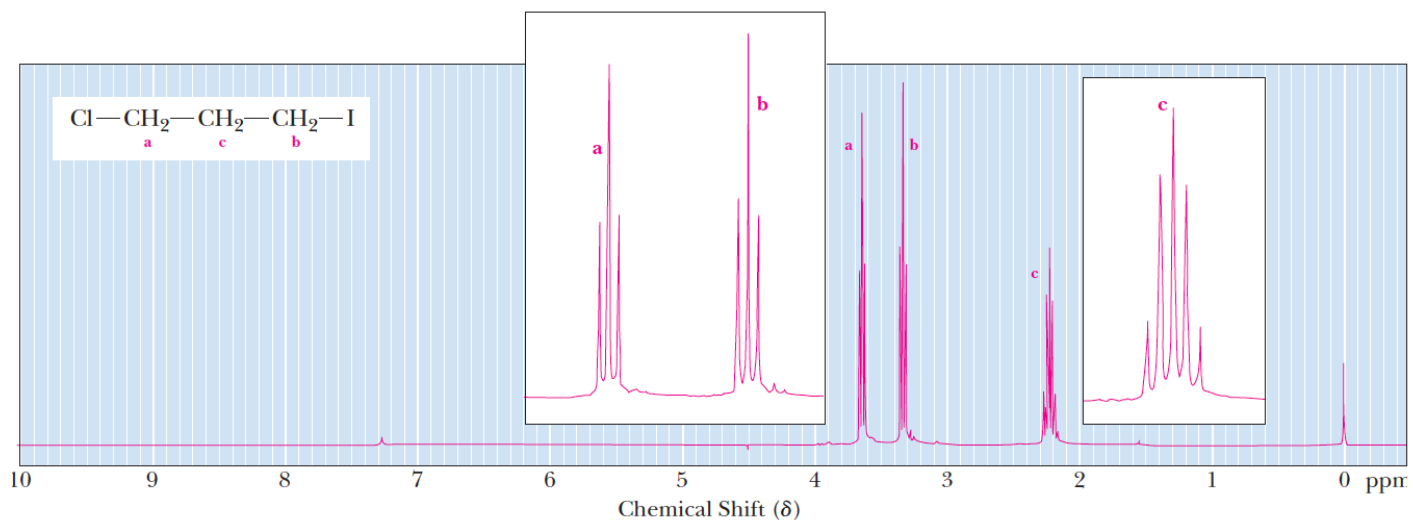


Figure 13.26

300 MHz $^1\text{H-NMR}$ spectrum of 1-chloro-3-iodopropane

M. For alkenes or ring structures such as cyclopropanes the splitting does not simplify (no bond rotation) and you see full multiplicative splitting ("doublet of doublets", etc.) [Click here to go to Pictures of the Day for today in which the NMR spectra for an alkene and a cyclic structure are explained.](#) Geminal coupling can be important for rings and alkenes.

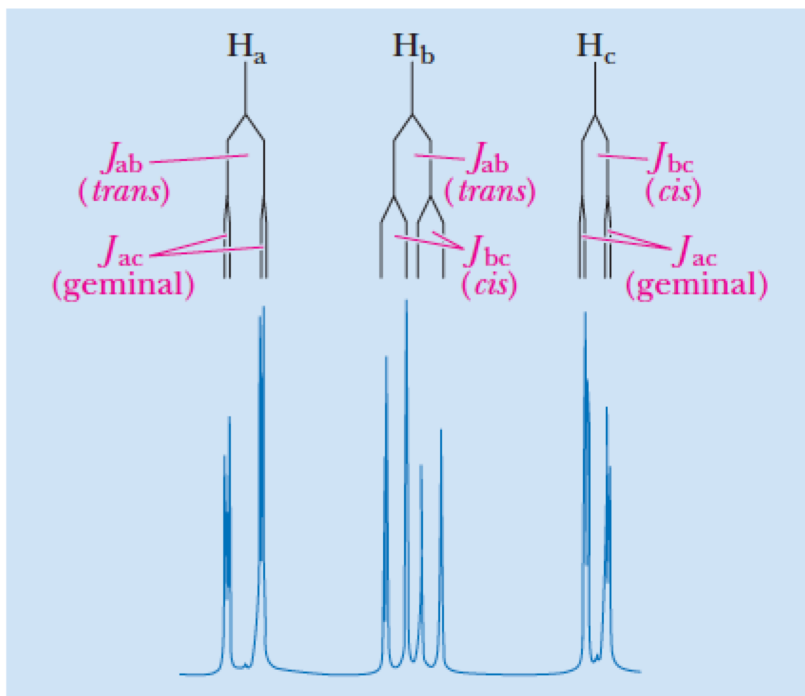
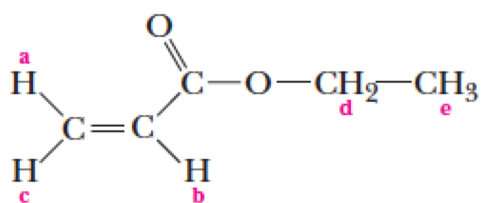


Figure 13.22

Tree diagrams for the complex coupling seen for the alkene H atoms in the $^1\text{H-NMR}$ spectrum of ethyl propenoate.

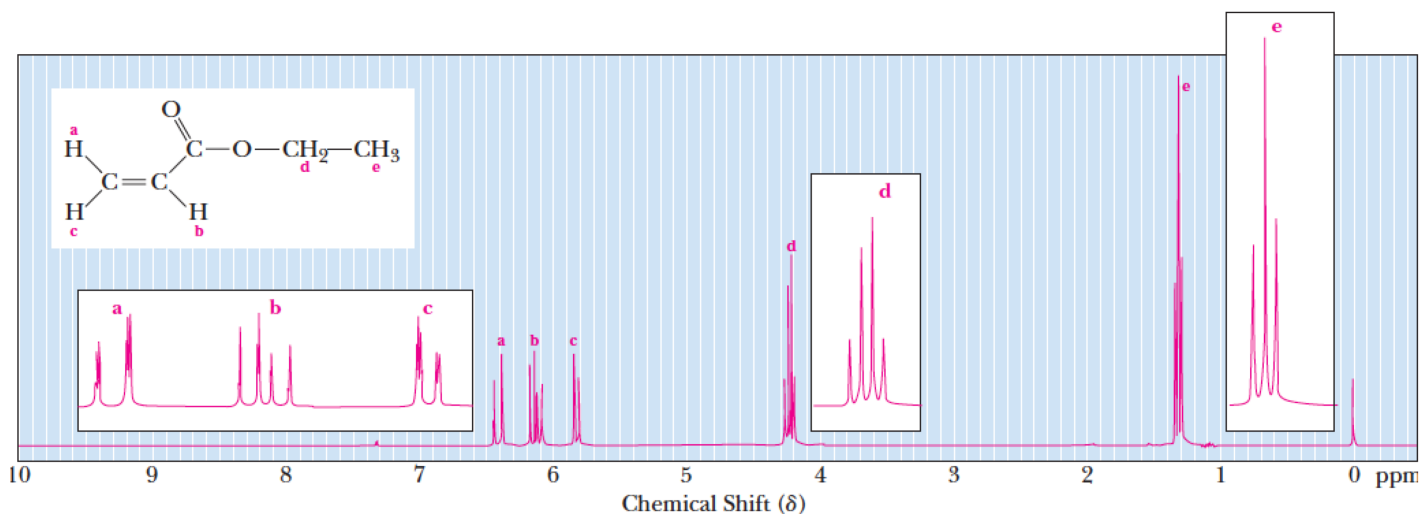


Figure 13.21

300 MHz $^1\text{H-NMR}$ spectrum of ethyl propenoate.

The popular medical diagnostic technique of **magnetic resonance imaging (MRI)** is based on **the same principles as NMR**, namely the **flipping (i.e. resonance) of nuclear spins of H atoms by radio frequency irradiation** when a patient is placed in **a strong magnetic field**. **Magnetic field gradients are used to gain imaging information**, and rotation of the gradient around the center of the object gives imaging in an entire plane (i.e. slice inside patient). In an MRI image, you are looking at **individual slices that when stacked make up the three-dimensional image of relative amounts of H atoms**, especially the H atoms from **water and fat, in the different tissues** [Memorize the preceding passage, as it will be worth 14 points on the final. No I am not kidding, 14 points right there.]