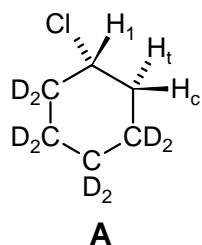
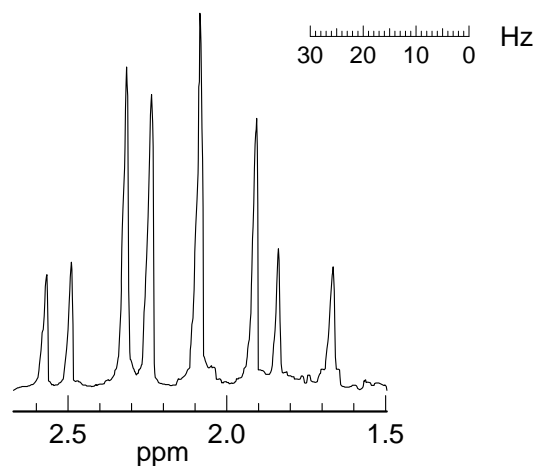
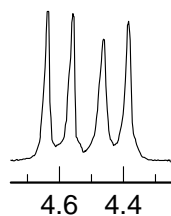


**Problem N-373** ( $C_6H_{11}Cl$ ). The deuterium-decoupled 60 MHz  $^1H$  NMR spectrum of **A** at  $-12.6\text{ }^\circ C$  is shown below. Clearly assign peaks to  $H_1$ ,  $H_c$  and  $H_t$ , and mark the distances corresponding to J values assuming an AMX analysis of the spectrum. Briefly tell how you made the assignments (digitized hard copy spectra from Hoefner, Lesko, Binsch *Org. Magn. Reson.* **1978**, 11, 179).

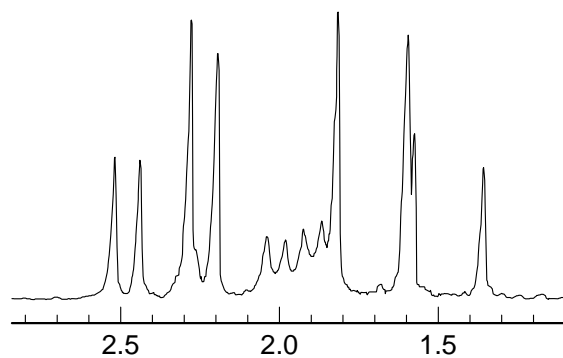
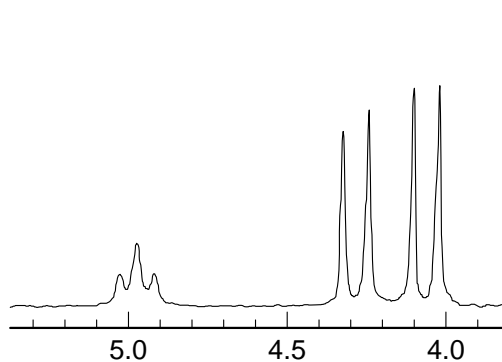


$T = -12.5\text{ }^\circ C$



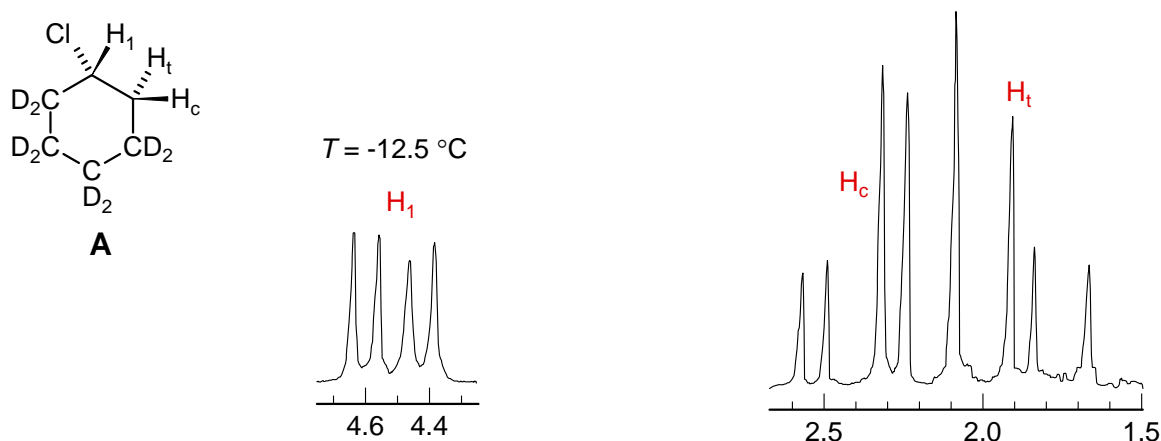
The low-temperature NMR spectrum of **A** is shown below. Clearly indicate assignments and explain the multiplicities observed.

$T = -76.2\text{ }^\circ C$



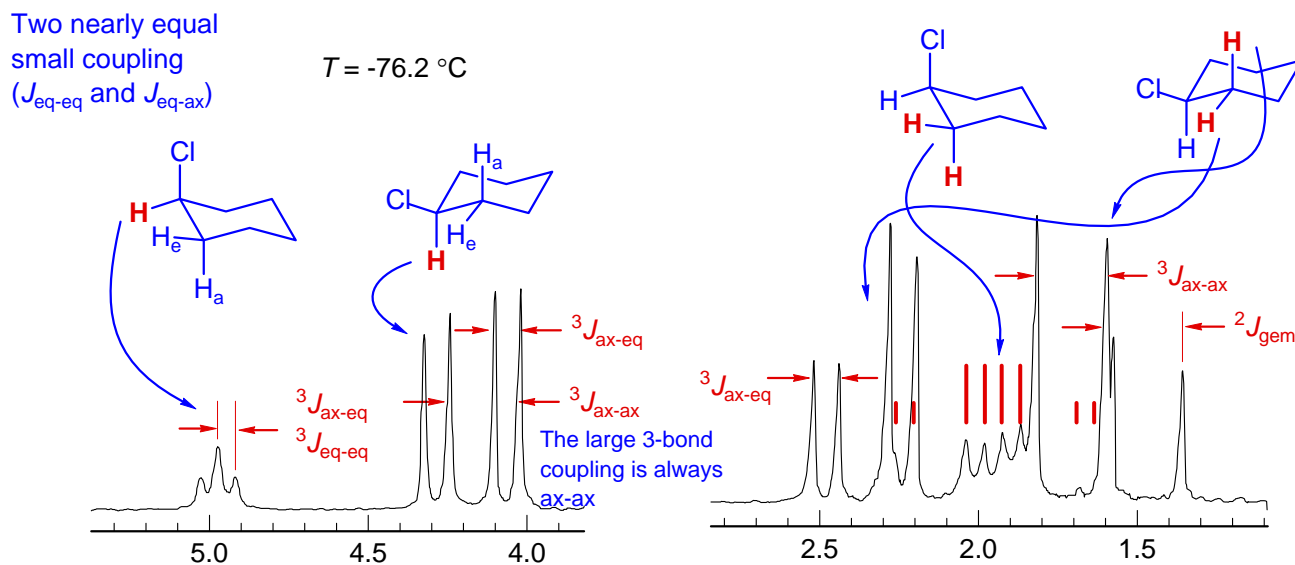
NOTE: Chemical shift scale may not be exact.

**Problem N-373** ( $C_6H_{11}Cl$ ). The deuterium-decoupled 60 MHz  $^1H$  NMR spectrum of **A** at  $-12.6\text{ }^\circ\text{C}$  is shown below. Clearly assign peaks to  $H_1$ ,  $H_c$  and  $H_t$ , and mark the distances corresponding to J values assuming an AMX analysis of the spectrum. Briefly tell how you made the assignments (digitized hard copy spectra from Hoefner, Lesko, Binsch *Org. Magn. Reson.* **1978**, 11, 179).



At  $-12.6\text{ }^\circ\text{C}$  ring inversion of chlorocyclohexane is fast on the NMR time scale, so one set of averaged signals is seen.

The low-temperature NMR spectrum of **A** is shown below. Clearly indicate assignments and explain the multiplicities observed.



At  $-76\text{ }^\circ\text{C}$  the ring inversion of chlorocyclohexane has become slow on the NMR time scale, so distinct signals are seen for the axial and equatorial conformations, with the characteristic small  $^3J_{eq-eq}$  and  $^3J_{ax-eq}$ , and the large  $^3J_{ax-ax}$ .