

THE INVERSION SPLITTING OF $^{15}\text{NH}_2\text{D}$ AND $^{15}\text{ND}_2\text{H}$ AS OBTAINED FROM THEIR FIR SPECTRA

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The far infrared absorption spectra of the partially deuterated species $^{15}\text{NH}_2\text{D}$ and $^{15}\text{ND}_2\text{H}$ have been recorded in the 20 to 105 cm^{-1} region using a Bruker IFS 120 Fourier transform spectrometer. A 25 cm long absorption cell with teflon windows was filled with different mixtures of $^{15}\text{NH}_3$ and $^{15}\text{ND}_3$ at total pressures ranging from 0.8 to 2 mbar. Several spectra with different $^{15}\text{NH}_2\text{D}$ to $^{15}\text{ND}_2\text{H}$ ratio were thus recorded. For all spectra the maximum path length was used resulting in a $2 \times 10^{-3} \text{ cm}^{-1}$ resolution.

Prior to the line assignment, rotational constants for $^{15}\text{NH}_2\text{D}$ and $^{15}\text{ND}_2\text{H}$ were calculated from those of $^{14}\text{NH}_2\text{D}$ and $^{14}\text{ND}_2\text{H}$. The IAM-type theoretical approach developed by Cohen and Pickett,^a which accounts for the large amplitude inversion motion and its coupling with the overall rotation, was afterwards used to obtain rovibrational energies and predicted spectra. In the case of $^{15}\text{NH}_2\text{D}$, observed line positions for low- J lines were sometimes within 0.050 cm^{-1} from their predicted values.

The line position analyzes are still in progress. For both species, they are carried out with the help of the IAM-type theoretical approach.^a For $^{15}\text{NH}_2\text{D}$ more than 300 a - and c -type lines have already been assigned and the observed wavenumbers are reproduced with an RMS deviation of $0.4 \times 10^{-3} \text{ cm}^{-1}$. This will be discussed in the paper and the values obtained for the spectroscopic constants will be reported and compared to those obtained by Fusina *et al.*^b for the ^{14}N -species. The decrease of the tunneling inversion splitting which goes from 0.406 cm^{-1} in $^{14}\text{NH}_2\text{D}$ ^b to 0.383 cm^{-1} in $^{15}\text{NH}_2\text{D}$ will also be discussed.

^aE. A. Cohen and H. M. Pickett, *J. Molec. Spectrosc.* **93**, 83 (1982).

^bL. Fusina, G. Di Lonardo, J. W. C. Johns and L. Halonen, *J. Molec. Spectrosc.* **127**, 240 (1988).