## MOLECULAR BEAM INFRARED SPECTRUM OF THE HF STRETCHING FUNDAMENTAL BAND OF THE HF-BF $_3$ COMPLEX

## K. NAUTA and R.E. MILLER, *Dept. of Chemistry, University of North Carolina, Chapel Hill, NC* 27599; G.T. FRASER and W.J. LAFFERTY, *Optical Technology Division, NIST, Gaithersburg, MD* 20899.

The microwave spectrum of the antihydrogen bonded HF-BF<sub>3</sub> complex has recently been reported by Philips et al. <sup>*a*</sup>. The spectrum reveals a near symmetric F-BF<sub>3</sub> heavy atom frame with the proton slightly off axis, and a small barrier to internal rotation of the H atom against the F atoms. Here, we have investigated the H-F stretching fundamental band of this complex using a molecular beam optothermal spectrometer and a color-center laser. Two subbands are expected: a parallel band with  $\Delta m = 0$  selection rules and a perpendicular band with  $\Delta m = \pm 1$  selection rules. At present, only the perpendicular band has been observed. This band has the appearance of a perpendicular band of a symmetric rotor, but with anomalously strong <sup>*r*</sup>R transitions and weak <sup>*p*</sup>P transitions. Upper state constants are obtained fixing the ground-state constants to the microwave-determined values. The band has a large positive  $\zeta$  Coriolis constant, from which an estimate of the barrier to internal rotation can be made. A search is presently underway to locate the expected weaker  $\Delta m=0$  parallel component.

<sup>&</sup>lt;sup>a</sup>J. A. Philips, M. Canagaratna, H. Goodfriend, A. Grushow, J. Almlof and K. R. Leopold J. Am. Chem. Soc. <u>117</u>, 12459 (1995)