INFRARED ABSORPTIONS OF THE H₂O—H₂ COMPLEX TRAPPED IN SOLID NEON

DANIEL FORNEY, <u>MARILYN E. JACOX</u>, AND WARREN E. THOMPSON, Optical Technology Division, National Institute of Standards and Technology, Gaithersburg, MD 20899-8441.

The infrared spectrum of a mixture of H_2O and H_2 with a large excess of neon which has been deposited at 4.2 K includes several new absorptions which can be assigned to the H_2O — H_2 van der Waals complex. The H–H stretching vibration of the H_2 moiety is detectable even for very dilute concentrations of the complex. Small shifts in the position of the OH-stretching absorptions associated with vibrotational transitions of the H_2O moiety also are observed. The intensities of absorptions arising from nonrotating H_2O are considerably enhanced for the complex. Studies of samples enriched in D_2O and in D_2 give similar results. Since HD, unlike H_2 and D_2 , is readily thermally depopulated to its J = 0 rotational state, this observation is consistent with the stabilization principally of the complex between H_2O and H_2 or D_2 in its J = 1 rotational state.