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solutions by analyzing the difference of the amplitudes of oscillatory water structure signals in the presence and absence of D₂O. We present a diffusion assay to measure and confirm diffusion in water-soluble proteins, by exploiting the difference of the amplitudes of oscillatory water structure signals in the presence and absence of deuterium oxide. The method is based on the change of the oscillation frequency of 2H(2)O structured water by the addition of D(2)O, which gives a peak shift, or a peak flattening, in the NMR spectrum. In this paper, the diffusion coefficients in lysozyme and a synthetic polymer were measured, and were shown to be consistent with the published values for these materials. The difference between the diffusion coefficient of the protein sample in D(2)O and in H(2)O also provides the amount of water bound to the protein. We then applied the method to protein G, a recombinant filovirus glycoprotein that possesses viral agglutination activity, and found that one molecule of the protein can bind one molecule of water.

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