

Albert Freedman, PhD, is a psychologist in independent practice, serving clients and families nationally. Dr. Freedman provides counseling support for patients and families affected by rare disease. He serves as a consulting psychologist to rare disease advocacy organizations and biopharmaceutical companies globally. Dr. Freedman speaks widely on the topic of mental health and rare disease. Al's son, Jack, lived with Spinal Muscular Atrophy (SMA) for 26 years.