dataMSA Surveys for Multiple System Atrophy: Patient Data Report

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When I couldn't find patient-driven information about MSA, and when my doctors didn't have a lot of information about patient experiences, I set out to create a way patients could share their experiences.dataMSA has collected information from nearly 350 persons worldwide who have been affected by MSA (patients, caregivers, loved ones to those lost). The data has been shared withresearchers at UCSF, Vanderbilt, Mayo Clinic, UF and other centers. This text is an invaluable resource for health care providers, support staff members and others who seek information about multiple system atrophy.No revenue is collected from the surveys and patients remain confidential. 100% of the profits from this book will be donated to the MSA Coalition to support research for a cure.Thank you for helping us raise our MSA voices. Bill Sydnor, dataMSA creator

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