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The Amyloidosis Foundation is a non-profit patient advocacy organization founded for, and by patients and families affected by this uncommon "orphan" disease. We are US based and organized exclusively for charitable purposes under Section 501(c)(3) of the Internal Revenue Code. Our core mission is **medical community awareness** of amyloidosis and its devastating effects. Why is this so important? Because we believe awareness will increase the rate of early detection, and lead to **improved survivability and quality of life**. We also provide for **patient education** and contribute to **research opportunities**. We sponsor programs and services, worldwide, through the leadership of our knowledgeable and dedicated volunteers. Our volunteers receive no compensation. Donations are distributed across Awareness, Patient Education and Research.

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