

We want to hear your ATS story!

We are connecting with you because you know someone who has arterial tortuosity syndrome (ATS). Because this disease is so rare, a better understanding of ATS is essential to create global awareness.

Cambridge BioMarketing, a global healthcare communications agency focusing on rare diseases, is currently in the process of creating an informational website for families affected by ATS. Because there are so few people with this disease, it can be hard to know where to start looking for materials. This website will be a centralized place to find information about ATS as well as the current ATS research.

In an effort to learn more about what it is like to have a loved one facing ATS, Cambridge BioMarketing is looking to interview the hardworking caregivers of people with ATS. The goal is to gain a better understanding of what life is like for people with ATS.

If you or someone in your family is interested in helping further society's understanding of ATS, we are eager to talk with you. You could share your experiences through a phone interview or via email.

If you would like to participate in helping us create this informational website and you consent to an interview, please contact Lin Betancourt or Julia Bond at your convenience using the information below.

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We look forward to hearing from you soon.

All the best,

Cambridge BioMarketing