

What Differentiates the TS Alliance

The TS Alliance is a model nonprofit in the rare disease research and support sector. Here are some ways we have demonstrated our unique ability to reach our constituents and impact their quality of life.

Provided support by a TS Alliance advocate to **4,083** individuals with TSC and facilitated **24,852** peer-to-peer connections in 2018, helping to reduce the stress and anxiety of a TSC diagnosis.

Established and built the first TSC Natural History Database as well as a TSC Biosample Repository and brought together a consortium of researchers who are now conducting the first preventative clinical trial in both TSC and epilepsy.

Raised
12.4
million
from more than 600 engaged donors and community members since launching the **Unlock the Cure** research funding campaign in 2011, thereby advancing TSC research.

Reached more than
21
million
people through the **#IAMTSC** awareness campaign, dramatically increasing the visibility of TSC.



Galvanized the TSC community, and through their advocacy efforts, the Department of Defense Congressionally Directed Medical Research Program has appropriated **\$83 million** toward TSC research since 2002.

Grown our volunteer base from **95** to more than **1,800** volunteers today, highlighting our community-building expertise.