Thank You for Your Continued Support!

Thank you for your donations and continued support of cureCADASIL! Your generous donations have allowed cureCADASIL to grow and continue to support CADASIL families and research through the years.

During 2018 the cureCADASIL team focused on:
- Creating 3 new CADASIL Connection webinars focused on research
- Creating and launching a new, patient-centered CADASIL care webcast series, focused on issues of importance to CADASIL families
- Recruiting CADASIL patients via the CADASIL Family Registry and social media for 2 prospective studies at UC San Francisco and Mayo Clinic Jacksonville
- Growing the cureCADASIL Family Registry for future CADASIL studies
- Supporting the Research Alliance for CADASIL research and expanding our research funding to other US researchers
- Upgrading our website to include research studies CADASIL patients can join, research cureCADASIL has supported, as well as all the recorded webinars and an online submission portal to submit patient/family stories
- Attending conferences to promote CADASIL research and make scientific connections

We wish all families with CADASIL and those that care for them all the best in the new year!

Looking back on accomplishments of 2018 as we get ready to Embrace the Challenge for 2019

Rare Disease Day is February 23! We are asking members of the CADASIL community to contribute 1-3 words that come to mind when thinking of CADASIL. Together we will create a community word art piece for Rare Disease Day like this one from last year.

We need you! Team CADASIL has been selected to participate in the 2019 Million Dollar Bike Ride to raise money for a CADASIL research grant to support through UPenn! If you are interested in joining Team CADASIL as a rider registration is now open!

CADASIL care webcasts began in 2018, focusing on topics important to patients and families based on surveys of the community.
The second webcast by researcher and clinician, Dr. Swati Sathe a member of the cureCADASIL Scientific Advisory Board, focuses on CADASIL and migraines and is now available on our website!

Karla's CADASIL story from Brazil

Karla is a 42-year-old administrative manager of a luxury residential condominium complex in Brazil who has been living with CADASIL for almost two decades. It took her doctors many years and many incorrect diagnosis to find this rare disease. Read more on our website.