



A better world for people with CHARGE syndrome

The CHARGE Syndrome Foundation champions the lifelong potential of people with CHARGE syndrome through outreach, education and research.

Contact Us For More Info

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Who are we?

The CHARGE Syndrome Foundation, Inc. is a 501(c)(3) charitable organization that is governed by a Board of Directors, guided by a group of Advisors, fueled by volunteers and staffed by a small team of dedicated professionals. The primary goals of the Foundation include: growing our outreach and support programs; increasing our knowledge base; promoting awareness of CHARGE syndrome; building networks and partnerships; and investing in leading-edge research.

Our Services



Family Assistance Programs

Medical Travel Program reimburses families for costs associated with travel to and from long-distance medical appointments

Ethan Wolfe Recreational Assistance Program (EWRAP) awards individuals with CHARGE adaptive recreational equipment and activities - because everyone deserves the chance to play!

Crocker Barker Sibling Scholarship awards college scholarships to siblings of individuals with CHARGE



Outreach

We provide support to the CHARGE community through our dedicated Outreach Support Manager, Family Liaisons, support groups, biennial international conferences, informational webinars, social media support groups, and newsletters

Research

Thanks to the generosity of our donors, over the years the Foundation has been able to fund over 1.3 million to CHARGE syndrome-related research projects. Our research grant recipients have gone on to publish their research in prestigious journals and receive close to \$12 million from the NIH and other government agencies.

Conference

At our biennial conferences, families get to share experiences, learn from one another, make lifelong friendships, experience a sense of belonging unlike any other, and hear from the world's top experts in CHARGE syndrome.

