

The FANNS (Family Advancement Network Services) Project was created to provide a database for families, where they can find resources to help them make informed decisions when navigating their child's health.



Our Global Delegates are those who are also making this rare disease journey. They have graciously accepted the responsibility of being a point of contact in their state or country and are ready to share their experiences. The most common advice they give, is to be connected with a VAC (Vascular Anomalies Clinic) Team.

The CMTC Alliance provides an annual conference, where families receive evidence-based & up-to-date informative presentations, by the leading experts in the country. Also available are clinical evaluations, free-of-charge, by our expert medical team.

In addition to the annual conference, a Family Event is planned the day before the conference to allow everyone the opportunity to get to know one another. This is an excellent time for the kids (of all ages) to play and realize they are not making this journey alone!

Moreover, we have an active closed group on Facebook, giving patients, parents, and care givers the freedom to ask questions, and sometimes vent when the journey follows an unfortunate path.



If you have any questions, please contact our president at: president@cmtcalliance.org

CMTC Alliance

3715 Wesley Chapel Rd
Zanesville, Ohio 43701
740-408-3537
<https://www.cmtcalliance.org>

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