



**ahcf**  
alternating hemiplegia of childhood foundation

CLINICAL REGISTRY

## AHC Clinical Registry

We encourage all AHC patients and their families to enroll in the AHC Registry.

The AHC Registry provides a mechanism to uniformly collect meaningful clinical data in an effort to enhance our understanding of the variability of AHC and other ATP1A3 related disorders. Collection of robust natural history data is important for us as a community to better understand AHC to help promote earlier diagnosis, better medical management and facilitate robust data to support future clinical trials.

We hope this collection of clinical data and associated samples from affected patients and their families will foster additional research collaborations that will ultimately help promote earlier diagnosis, increase knowledge about challenges in medical management and provide a means to engage the broadest possible participation in clinical studies and trials so critical to promote progress in finding effective treatments for this disorder.

For questions regarding enrollment in the AHCF registry please contact:

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