To ALL SCN2A Families,

*The time to make your voices heard for SCN2A research is here! We encourage you to participate in this study that we have developed to prepare us for clinical trials coming in the near future. Every patient, **regardless of age**, with a disease-causing SCN2A variant should register and go through the initial survey **even if there is no epilepsy diagnosis**. This is a first step. We will not stop until every SCN2A related disorder is cured.*

—The FamilieSCN2A Foundation Executive Director and Board of Directors

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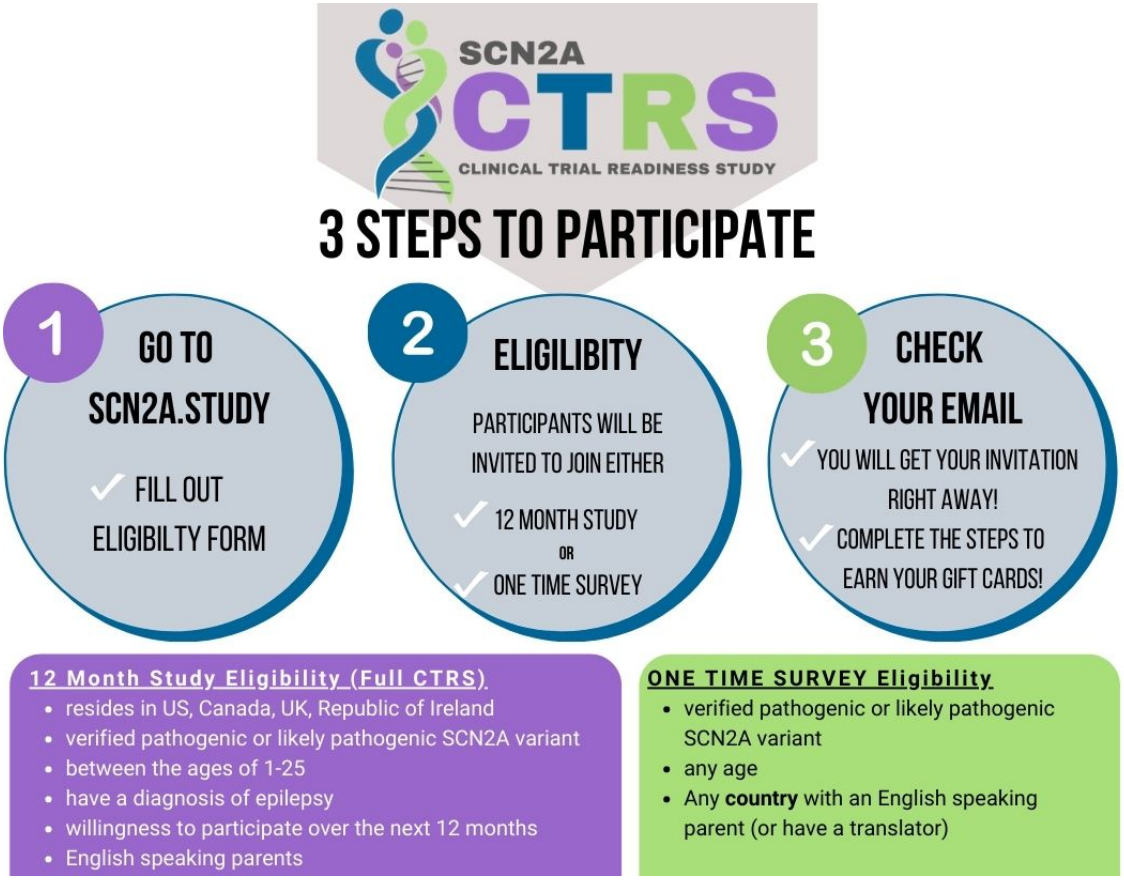

Click here for more info and registration

Please contact us at impact@scn2a.org with any questions. As always, you can learn about research opportunities on our website under the RESEARCH tab.

SCN2A RELATED AUTISM & EPILEPSY
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