I ❤️ SOMEONE WITH SLC13A5 EPILEPSY

TESS RESEARCH FOUNDATION for SLC13A5 Epilepsy
tessresearch.org

#SLC13A5Awareness #TESSResearch
Five Things You Can Do To Spread Awareness of SLC13A5 Epilepsy

1. Get Social with this Sign and spread #SLC13A5Awareness

2. Raise Money for SLC13A5 Research and donate to #TESSResearch.

3. Wear purple to show your SLC13A5 Epilepsy pride.

4. Watch and share our new video at tessresearch.org.

5. Educate a friend about SLC13A5 Epilepsy using these five things to know.

Five Things To Know About SLC13A5 Epilepsy

SLC13A5 Epilepsy is a rare form of genetic epilepsy with seizures beginning within the first days of life.

People affected by SLC13A5 Epilepsy require 24-hour care. They have trouble walking, speaking, sleeping, and eating.

SLC13A5 Epilepsy knows no geographic boundaries. TESS helps families around the world.

There are currently no treatments to cure this disease.

TESS Research Foundation, a 501(c)(3) non-profit, is the only organization in the world dedicated to supporting families and driving research to treat and cure SLC13A5 Epilepsy.