

I  **SOMEONE WITH  
SLC13A5 EPILEPSY**

 **TESS** **RESEARCH**  
FOUNDATION  
for SLC13A5 Epilepsy

[tessresearch.org](https://tessresearch.org)

**#SLC13A5Awareness**

**#TESSResearch**

# Celebrate SLC13A5 Epilepsy International Awareness Day May 13th



[tessresearch.org](http://tessresearch.org)

## 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](http://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.

[tessresearch.org](http://tessresearch.org)