



Glut1 Deficiency Foundation
PO Box 737 Owingsville, KY 40360
Glenna Steele, Executive Director
gsteele@G1DFoundation.org

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To Whom It May Concern,

We are writing in support of the Maryland Seizure Safe School Act, or Brynleigh Act, which calls for state legislation to ensure all school personnel, including nurses, teachers, and volunteers, are not only prepared but can recognize and respond appropriately and efficiently to students experiencing seizures. Approximately 1 in 26 people will develop epilepsy in their lifetime, 1 in 10 people will have a single seizure in their lifetime, and approximately 7900 Maryland children currently have an active epilepsy diagnosis, but there is only one nurse per 913 students. It is of vital importance that school staff know how to recognize and respond to a seizure should one occur in their classroom, school bus, or specialized area. Quick response to a seizure is vital, and cannot wait for a nurse to arrive. It is not enough for school nurses to be the only ones with that training. Many students will have their first seizure while in school, so this is not just a matter of following a plan that is already in place.

The Brynleigh Act will ensure that all school personnel have some familiarity with seizure recognition, and in doing so, will make all our children safer in their school environment. Four other states have passed similar bills, and many other states will follow. We hope that Maryland will, as always, lead by example. We stand with the Epilepsy Foundation in supporting this legislation.

The Glut1 Deficiency Foundation is a nonprofit family organization dedicated to improving the lives of those in the Glut1 Deficiency community through its mission of increased awareness, improved education, advocacy for patients and families, and support and funding for research. Glut1 Deficiency is a rare genetic disorder of brain metabolism, and the vast majority of our patients have seizures of varying types and frequency.

Sincerely,

Glenna Steele
Executive Director
Glut1 Deficiency Foundation
gsteele@G1DFoundation.org
859-585-2538