MOG Antibody Disease Fact Sheet for Educators

Accommodations

Considerations

Learning

Classroom

- MOG Antibody Disease (MOG-AD) is a rare neuroimmune condition that typically causes inflammation in the optic nerve, spinal cord and/or brain.
- MOG-AD is an immune disease, which occurs, in at least part, due to attack of a person's own immune cells on the MOG protein present in their brain, optic nerve or spine.
- MOG-AD is not infectious and cannot be passed on to others (i.e., it is not transmissible)
- There is no cure, but treatments are available to prevent inflammatory attacks and to manage symptoms.
- MOG-AD symptoms often come (relapses) and go (remissions). Patients can feel well one day and can have new symptoms the next.
- MOG-AD relapses are not brought on by physical activity or by reading or schoolwork. However, during relapse, and as a child recovers from a relapse, schoolwork may need to be adjusted.
- Loss or blurring of vision in one or both eyes Seizures Loss of color vision Vomiting Eve pain Headaches Paralysis or weakness of a limb or limbs Altered mental status (requires emergency Loss of sensation care), defined as a loss of awareness, reduced Cognitive issues (e.g., learning, memorization, responsiveness, confusion or loss of concentration) consciousness. Altered mental status can Behavioral changes/issues occur suddenly (as in the case of seizure) or Loss of bladder or bowel control more gradually over a few days in a severe Fatigue related to the diagnosis or medications relapse.
 - Short-term treatments to reduce inflammation during an acute attack include IV or oral steroids, plasma exchange (PLEX), and intravenous immunoglobulin (IVIg).
 - Students diagnosed with MOG-AD may be on long-term treatment with medications such as intravenous (IVIg) or subcutaneous (SCIg) immunoglobulin or medications that suppress the immune system, such as mycophenolate mofetil (CellCept), rituximab (Rituxan), azathioprine (Imuran), or lowdose steroids.

Other Considerations

- Some treatments carry an increased risk of infection to the student with MOG-AD, so it is important to keep the classroom clean and sanitized.
- Good hygiene and hand washing are important.
- Alert parents/guardians to any illnesses in the classroom (e.g., flu, strep throat, stomach virus).
- An action plan for medical emergencies, including seizure should be put in place.
- Provide accommodations, required by law, for students who use wheelchairs or other mobility or assistive devices.
- An emergency plan should be in place for exiting the building, medical emergencies, and a seizure plan (when appropriate).
- Student may need plans in place to assist with learning challenges (e.g., 504, IEP, EHC).
- Be cognizant of potential vision issues and their impact on learning.
- Multiple absences are common due to doctor appointments, multi-hour infusions, MRIs, and adverse treatment reactions. Providing a "homework buddy" to ensure that the MOG-AD patient keeps pace with school assignments, as well as taping key lectures for later learning are helpful.
- Inform parents/guardians of any changes in behavior (e.g., anger outbursts, anxiety, crying, student acting withdrawn) or new learning challenges.
- Student may be struggling with their diagnosis and the changes MOG-AD has caused in their life. School counselors may be very helpful.
- Consult with student's parents/guardians regarding privacy preferences around their condition.

Parents/caregivers should discuss the student's current, specific neurological symptoms with educators. Parents should be made aware of any new symptoms or anything out of the ordinary immediately, as symptoms vary widely, and some symptoms may require emergency care.

Note: Some symptoms may be triggered as a result of prolonged exposure to heat or over-exertion that leads to an increase in body temperature (Uhthoff's Phenomenon).



For more information on MOG Antibody Disease, please visit: https://wearesrna.org/living-with-myelitis/disease-information/mog-antibody-disease/

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Symptoms

Treatments