

Series 2:
Associated Symptoms

MOGbytes™

**A snack-sized guide to
understanding MOGAD**



a series brought to you by



The MOG Antibody: A Clue, Not the Whole Story



The MOG antibody is a biomarker found in blood or spinal fluid that is needed for doctors to diagnose MOGAD. Discovering the MOG antibody has helped doctors better understand and identify MOGAD.



False positives can happen; a positive MOG antibody test alone does not confirm MOGAD.



Doctors also look at symptoms, exam findings, and MRI results



Other conditions that can look like MOGAD, such as MS and NMOSD, must be ruled out

Will an Attack Happen Again?



MOGAD can follow two different patterns:



Monophasic

- One attack only, known as a single demyelinating event
- About half of all people with MOGAD may be monophasic

Doctors sometimes observe this pattern in people who:

- Had ADEM symptoms during their first attack
- May have MOG antibody levels that decrease over time



Relapsing

- More than one attack, known as multiple demyelinating events

Doctors sometimes observe this pattern in people who:

- First develop symptoms in adulthood
- Had symptoms other than ADEM during their first attack
- Continue to have positive MOG antibody tests

Currently, there is no reliable test to predict who will relapse.

Where Does MOGAD Affect the Body?



MOGAD can affect different parts of the central nervous system, such as:



Optic Nerve



Spinal Cord



Brian/Brainstem

These areas may be affected all at once or at different times.

How Does MOGAD Show Up in the Eye?



Inflammation of the Optic Nerve

The optic nerve is behind the eye. This is known as optic neuritis (ON) and can lead to:



Changes in vision, such as blurry vision, vision loss, or blindness



Eye Pain



Headache



How Does MOGAD Show Up in the Spinal Cord?



Inflammation of the Spinal Cord

This is known as myelitis or transverse myelitis (TM)
and can lead to:



**Muscle numbness,
weakness, pain, and
spasms**



**Paralysis below the
point of damage in the
spinal cord**



How Does MOGAD Show Up in the Brain?



Inflammation of the Brain, Brainstem, and/or Meninges

The meninges are the covering of the brain and spinal cord.

This can lead to:



Autoimmune Encephalitis (AE)

**Acute Disseminated
Encephalomyelitis (ADEM)**

Cerebral Cortical Encephalitis (CCE)

Brainstem Encephalitis

Meningitis

Symptoms of the above may include headaches, fever, confusion, seizures, loss of coordination and/or balance, loss of consciousness, brain fog, difficulty swallowing, and behavioral changes.

How Can MOGAD Differ by Age?



Some presentations are more common in MOGAD depending on age.



Children

- The most common presentation in children is ADEM
- Around 50% of patients with ADEM test positive for MOG antibodies
- In one study, MOG antibodies were the most commonly identified antibody in children with autoimmune encephalitis



Adults

- The most common presentation in adults is ON
- ON is also the most common initial symptom of MOGAD, seen in about 30–60% of adult patients

Because MOGAD can look different in each person, new or worsening symptoms may signal a relapse and should be discussed with a healthcare provider.

ADEM, acute disseminated encephalomyelitis; MOG, myelin oligodendrocyte glycoprotein; MOGAD, myelin oligodendrocyte glycoprotein antibody-associated disease; ON, optic neuritis

What Happens After an Attack?



After recovering from the first attack, doctors may **monitor closely** before starting long-term treatment.



After relapse attacks, doctors may **change the treatment plan**.



Every situation is different. Treatment decisions are personalized and should be made together with the patient and healthcare provider.



Understanding Relapses in MOGAD



Around half of all patients will experience a relapse.



Relapses are hard to predict, and they can affect the same or different areas.



ON is common in relapses, especially for those who have already experienced it.

Many people can recover from MOGAD relapses. However, repeated relapses can sometimes lead to lasting disability.

Looking Ahead



Researchers are still studying this disease as MOGAD was recognized as a distinct disease relatively recently.



Doctors are learning more every year to improve diagnosis, treatment, and long-term care.



Awareness is growing, and it is up to us advocate for ourselves and loved ones to receive a quicker and more accurate diagnosis.

Disclaimer:

If you are experiencing any of these neurological symptoms, ask your doctor whether you should be tested for the MOG antibody through a live cell-based assay MOG antibody test.

References



- Armangue, T., Olivé-Cirera, G., Martínez-Hernandez, E., Sepulveda, M., Ruiz-Garcia, R., Muñoz-Batista, M., Ariño, H., González-Álvarez, V., Felipe-Rucián, A., Jesús Martínez-González, M., Cantarín-Extremera, V., Concepción Miranda-Herrero, M., Monge-Galindo, L., Tomás-Vila, M., Miravet, E., Málaga, I., Arrambide, G., Auger, C., Tintoré, M., ... Villar-Vera, C. (2020). Associations of paediatric demyelinating and encephalitic syndromes with myelin oligodendrocyte glycoprotein antibodies: A multicentre observational study. *The Lancet Neurology*, 19(3), 234–246. [https://doi.org/10.1016/s1474-4422\(19\)30488-0](https://doi.org/10.1016/s1474-4422(19)30488-0)
- Gklinos, P., & Dobson, R. (2024). Myelin oligodendrocyte glycoprotein-antibody associated disease: An updated review of the clinical spectrum, pathogenetic mechanisms and therapeutic management. *Antibodies*, 13(2), 43. <https://doi.org/10.3390/antib13020043>
- Hooshmand, S. J., Vorasoot, N., Li, X., Chen, J. J., Redenbaugh, V., Banks, S. A., Cacciaguerra, L., Valencia-Sanchez, C., Lopez Chiriboga, S., Sechi, E., Tillema, J.-M., Tajfirouz, D., Basso, M., Hardy, T. A., Tisavipat, N., Guo, K., Pittock, S. J., & Flanagan, E. P. (2025). Application of the ADEM definition to cerebral attacks of MOG antibody-associated disease. *Multiple Sclerosis Journal*, 31(14), 1629–1640. <https://doi.org/10.1177/13524585251359450>
- Jain, K., Dhamija, K., Holla, V. V., Thennarasu, K., Bhat, M., Mahadevan, A., & Netravathi, M. (2025). MOGAD treatment outcomes and relapse characteristics: Real world experience from a Low-Middle income country. *Multiple Sclerosis and Related Disorders*, 100, 106548. <https://doi.org/10.1016/j.msard.2025.106548>
- Jeyakumar, N., Lerch, M., Dale, R. C., & Ramanathan, S. (2024). MOG antibody-associated optic neuritis. *Eye*, 38(12), 2289–2301. <https://doi.org/10.1038/s41433-024-03108-y>
- Li, Y., Liu, X., Wang, J., Pan, C., & Tang, Z. (2022). Clinical features and imaging findings of myelin oligodendrocyte glycoprotein-IgG-associated disorder (MOGAD). *Frontiers in Aging Neuroscience*, 14. <https://doi.org/10.3389/fnagi.2022.850743>
- Montalvo, M., Khattak, J. F., Redenbaugh, V., Britton, J., Sanchez, C. V., Datta, A., Tillema, J., Chen, J., McKeon, A., Pittock, S. J., Flanagan, E. P., & Dubey, D. (2022). Acute symptomatic seizures secondary to myelin oligodendrocyte glycoprotein antibody-associated disease. *Epilepsia*, 63(12), 3180–3191. <https://doi.org/10.1111/epi.17424>
- Santoro, J. D., Beukelman, T., Hemingway, C., Hokkanen, S. R. K., Tennigkeit, F., & Chitnis, T. (2023). Attack phenotypes and disease course in pediatric MOGAD. *Annals of Clinical and Translational Neurology*, 10(5), 672–685. <https://doi.org/10.1002/acn3.51759>
- Wang, J., Yang, K., Zhang, F., Yi, Y., & Wang, J. (2023). Clinical risk factors for recurrence of myelin oligodendrocyte glycoprotein antibody-associated disease. *Multiple Sclerosis and Related Disorders*, 77, 104879. <https://doi.org/10.1016/j.msard.2023.104879>

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Driven by **science.**

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