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JANUARY 2023

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FEATURED



Help Find MG Treatments through the MGFA Global MG Patient Registry

MGFA Global MG Patient Registry is a safe, secure, confidential means for MG patients to contribute actual patient data that will help researchers make MG discoveries that could lead to better treatments. Your contributions to the registry will help the entire MG Community. Register and add your data.

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TO BE MORE THAN
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Introducing New MGFA Board Member Callum Schjerner

MGFA welcomes our newest Board member Callum Schjerner. We are grateful for his expertise and commitment to the MG Community. Read more about Cal.

[Learn More»](#)

Seronegative MG Resource Center Now Open

Between 10 - 20% of people diagnosed with MG are seronegative. Patient information is scarce, but more is being done to research and develop better treatments for this “rare-of-the-rare” disease. Check out our Seronegative Resource Center. [Learn More»](#)



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Driven by science.

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Come Together at the MGFA National Patient Conference



Be Prepared in Advance for an MG Crisis

Living with MG can be difficult at times, and it is important to be



Hot new MG research targets MuSK MG with CAAR-T

Engineering T cells to target MuSK MG

The MG Community is gathering for the first time in three years! Join us in person in New Orleans on March 26-28.

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prepared for the future. Knowledge is key to understanding how to deal with a crisis, flare, or exacerbation.

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leads to clinical trial design. Read the latest research by Dr. Aimee Payne.

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Join the MGFA Online Community to Connect with Patients

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RESEARCH



Latest MG Research Clinical and Research Trials



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Is this monthly e-newsletter valuable for you and your family? Let us know. You can send comments directly to MGFA@myasthenia.org or contact Michael Antonellis at mantonellis@myasthenia.org.



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