

Accelerating myasthenia gravis diagnosis and treatment: Practical strategies and therapeutic advances

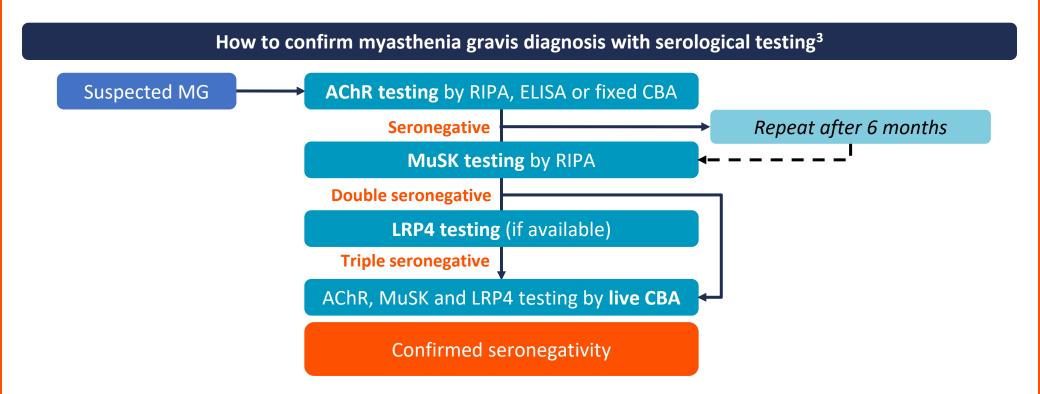
Practice aid for myasthenia gravis

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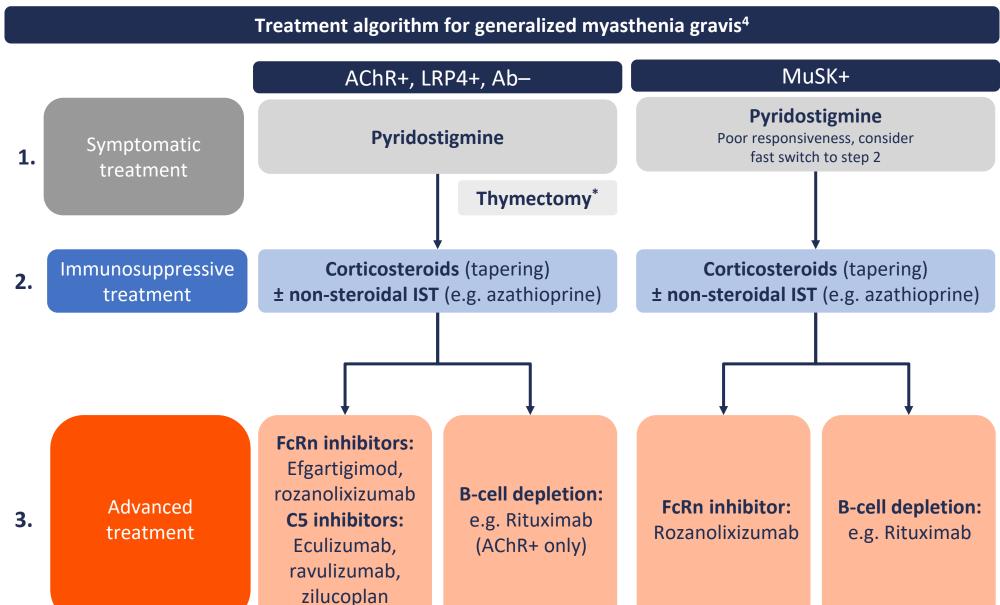
When to suspect myasthenia gravis

Most frequent presenting symptoms include^{1,2}

- Extraocular muscle weakness: Ptosis, double vision
- Bulbar muscle weakness: Difficulty chewing, choking, dysphagia
- Limb weakness: Upper limbs usually more impacted than lower limbs



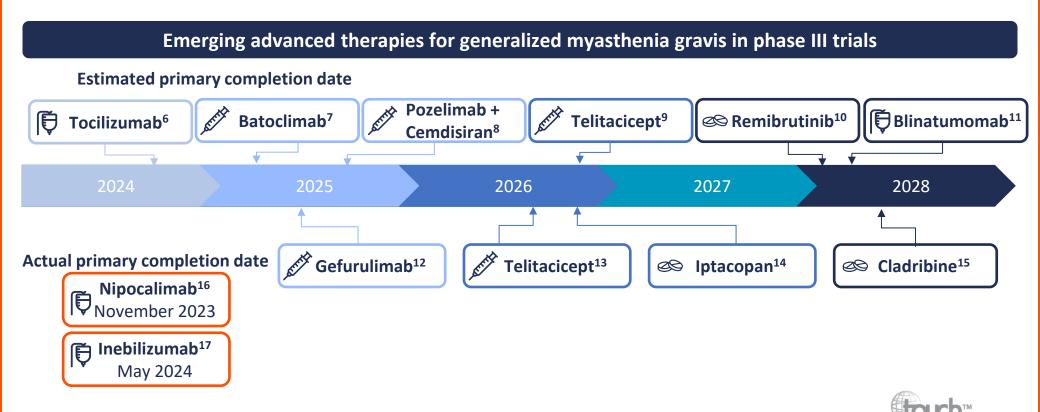






Approved advanced therapies for generalized myasthenia gravis⁵

Complement inhibitors	FcRn antagonists
<section-header> Eculizumab</section-header>	₱ ₱ Efgartigimod*
Ravulizumab	Rozanolixizumab
🞢 Zilucoplan	



^{*}A prefilled syringe formulation was approved on 10 April 2025 allowing for administration by patients and/or caregivers. 18

Abbreviations and references

Abbreviations

Ab, antibody; AChR, acetylcholine receptor; C5, C5 component of complement; CBA, cellular assays; ELISA, enzyme-linked immunosorbent assay; FcRn, neonatal Fc receptor; IST, immunosuppressive therapies; IV, intravenous, LRP4, low-density lipoprotein receptor-related protein 4; MG, myasthenia gravis; MuSK, muscle-specific kinase; RIPA, radioimmunoprecipitation assays.

References

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The guidance provided by this practice aid is not intended to directly influence patient care. Clinicians should always evaluate their patients' conditions and potential contraindications and review any relevant manufacturer product information or recommendations of other authorities prior to consideration of procedures, medications or other courses of diagnosis or therapy included here.

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