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## **Reducing burden of disease in gMG: Exploring the role of FcRn inhibitors**

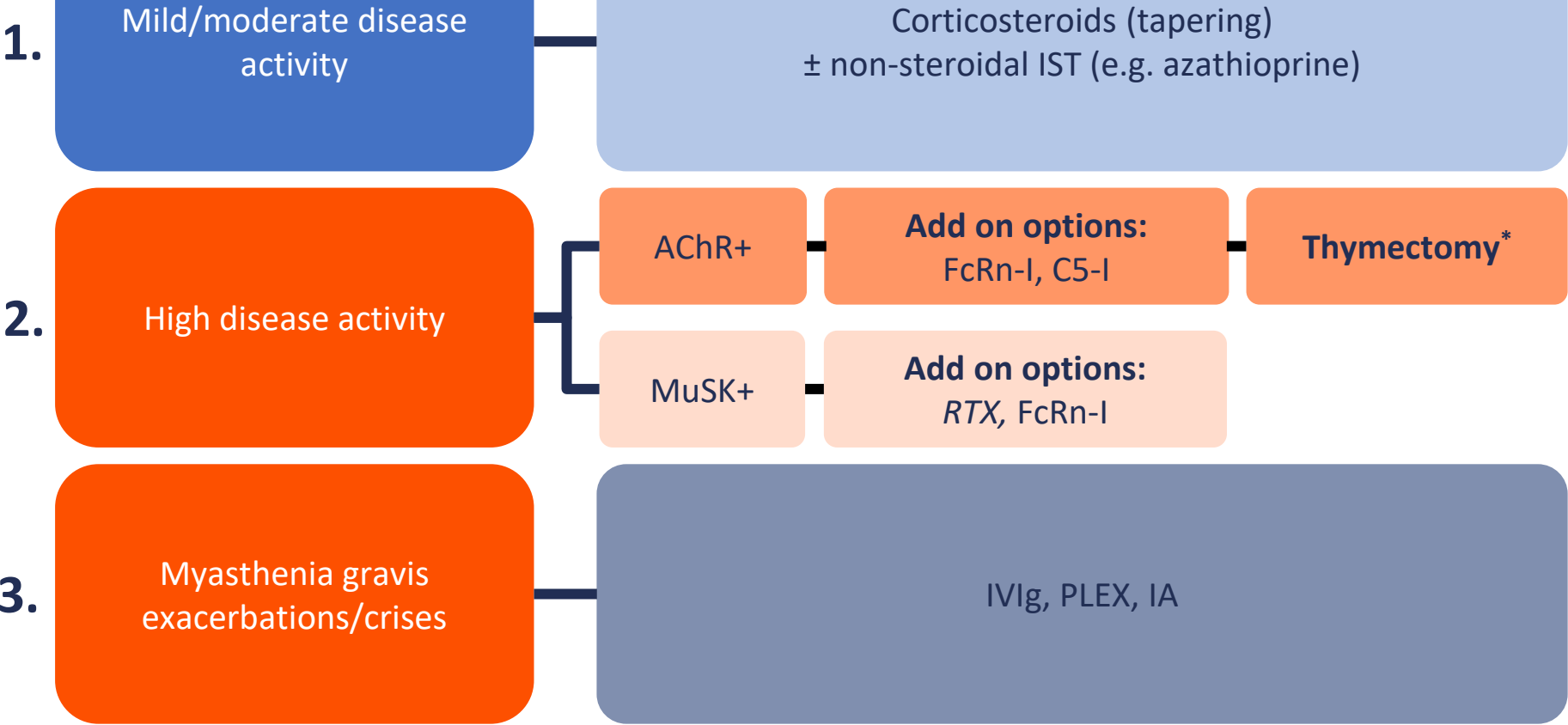
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**Practice aid for myasthenia gravis**

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Disease-modifying therapy options in generalized myasthenia gravis<sup>1,2</sup>

Symptomatic treatment with AChE inhibitors (e.g. pyridostigmine)



Treatments listed in italics are used off-label.  
\*Thymectomy is recommended in AChR+ patients who are eligible and should be performed within 2 years after diagnosis.

Approved FcRn inhibitor options in generalized myasthenia gravis<sup>3,4</sup>

	Efgartigimod	Rozanolixizumab
Dose	10 mg/kg (IV) 1,000 mg (SC)	7 mg/kg, 10 mg/kg, 15 mg/kg
Dosing regimen	Weekly IV for 4 weeks* <sup>†</sup> Weekly SC for 4 weeks* <sup>†</sup>	Weekly SC for 6 weeks*
Pharmacokinetic advantage	Rapid onset of action	SC administration convenient
US FDA and EMA approval status	Approved for use in AChR+ gMG	Approved in AChR+ and MuSK+ gMG

Emerging FcRn inhibitor options in generalized myasthenia gravis<sup>5</sup>

	Nipocalimab	Batoclimab
Dose	30 mg/kg initial dose followed by 15 mg/kg	680 mg
Dosing regimen	IV every 2 weeks*	Weekly SC for 6 weeks*
Pharmacokinetic advantage	Significant and sustained IgG reduction	SC administration convenient
US FDA and EMA approval status	Granted priority review by the FDA	N/A

\*Further continuation depending on treatment response; <sup>†</sup>Safety of starting subsequent cycles sooner than 50 days from the start of the previous cycle has not been established.<sup>3,4</sup>

## Factors to consider when managing generalized myasthenia gravis with FcRn inhibitors



Method of administration<sup>3,4</sup>



Type of generalized myasthenia gravis  
(AChR+ or MuSK+)<sup>3,4</sup>



Frequency of administration<sup>3,4</sup>



Concomitant use with other medications  
that also bind to FcRn<sup>3,4,7</sup>



Dosing schedule:  
cyclic vs predictable dosing<sup>6</sup>



Timing of vaccinations, particularly live or  
live-attenuated<sup>3,4</sup>



Time taken to administer dose<sup>3,4</sup>



Presence of active infections prior to  
treatment initiation<sup>3,4</sup>

## Abbreviations and references

### Abbreviations

ACh, acetylcholine; AChE, ACh esterase; AChR, ACh receptor; C5, C5 component of complement; EMA, European Medicines Agency; FcRn, neonatal Fc receptor; FDA, Food and drug administration; gMG, generalized myasthenia gravis; I, inhibitor; IA, immunoadsorption; Ig, immunoglobulin; IST, immunosuppressive therapy; IV, intravenous; MuSK, muscle-specific tyrosine kinase; N/A, not applicable; PLEX, plasma exchange; RTX, rituximab; SC subcutaneous.

### References

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3. FDA. Prescribing information. Available at: [www.accessdata.fda.gov/scripts/cder/daf/index.cfm](http://www.accessdata.fda.gov/scripts/cder/daf/index.cfm) (accessed 26 March 2025).
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5. Menon D, Bhandari V. *Expert Opin Emerg Drugs*. 2025; doi.org/10.1080/14728214.2025.2458061.
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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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