Prior Authorization Checklist for gMG Complement Inhibitors

A common reason for denial is incomplete or missing information on a prior authorization form. The following list provides standard information that payers typically request on a PA:

CLINICAL CRITERIA	MEDICATION AND VACCINATION HISTORY
Patient and healthcare prescriber information	Previous and current therapies, including name, duration of treatment, reason for
gMG ICD-10 Diagnosis Code (G70.00, G70.01)	discontinuation (if applicable):
MGFA Clinical Classification (Class II-IV)	Corticosteroids
☐ MG-ADL total score ≥6	immunosuppressive therapies
Positive serologic test for	 Immunosuppressive therapy (IVIg, PLEX, etc.) FcRn receptor antagonists
anti-AChR antibodies	Complement inhibitors
Signs and symptoms of gMG (date of onset, severity, exacerbations), comorbidities, etc.	Patient's meningococcal vaccinations, including dates of initial, second dose, and third doses, and boosters (if applicable):
	MenACWY
	MenB-4C or MenB-FHbp
	MenABCWY

Sign and date all necessary forms; the PA may be rejected if a signature is missing. Your office may need to coordinate with other providers to gather all necessary information to submit a PA.

*This checklist is provided as an educational resource only regarding common PA requirements for patients being prescribed a gMG complement inhibitor. Contact the individual payer for requirements and clinical coverage guidelines.

AChE=acetylcholinesterase; AChR=acetylcholine receptor; FcRn=neonatal Fc receptor; gMG=generalized myasthenia gravis; IVIg=intravenous immunoglobulin; ICD-10-CM, International Classification of Diseases, 10th Revision, Clinical Modification; MenACWY=quadrivalent (serogroups A, C, W, and Y) meningococcal conjugate vaccine; MenB=serogroup B meningococcal vaccines; MenB-4C=4-component meningococcal group B; MenB-FHbp=meningococcal serogroup B factor H binding protein; MG-ADL=Myasthenia Gravis-Activities of Daily Living; MGFA= Myasthenia Gravis Foundation of America; PLEX=plasma exchange; PA=prior authorization

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Prior Authorization Checklist for gMG FcRn Receptor Antagonists

A common reason for denial is incomplete or missing information on a prior authorization form. The following list provides standard information that payers typically request on a PA:

CLINICAL CRITERIA	MEDICATION HISTORY
Patient and healthcare prescriber information	Previous and current therapies, including name, duration of treatment reason for
gMG ICD-10 Diagnosis Code	treatment, reason for discontinuation (if applicable):
(G70.00, G70.01)	AChE inhibitors
MGFA Clinical Classification	Corticosteroids
(Class II-IV) \square MG-ADL total score ≥ 3	Non-steroidal immunosuppressive therapies
	Immunosuppressive therapy (IVIg, PLEX, etc.)
Positive serologic test for anti-AChR antibodies or	EcRn receptor antagonists
anti-MuSK antibodies	Complement inhibitors
Signs and symptoms of gMG (date of onset, severity, exacerbations), comorbidities, etc.	

Reminder: Some PA request forms may also require you to indicate the intended site of care for FcRn administration. If applicable, ensure the intended site of care is captured.

Sign and date all necessary forms; the PA may be rejected if a signature is missing. Your office may need to coordinate with other providers to gather all necessary information to submit a PA.

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AChE=acetylcholinesterase; AChR=acetylcholine receptor; FcRn=neonatal Fc receptor; gMG= generalized myasthenia gravis; IVIg=intravenous immunoglobulin; ICD-10-CM, International Classification of Diseases, 10th Revision, Clinical Modification; MenACWY=quadrivalent (serogroups A, C, W, and Y) meningococcal conjugate vaccine; MenB=serogroup B meningococcal vaccines; MenB-4C=4-component meningococcal group B; MenB-FHbp=meningococcal serogroup B factor H binding protein; MG-ADL=Myasthenia Gravis-Activities of Daily Living; MGFA= Myasthenia Gravis Foundation of America; MuSK=muscle-specific tyrosine kinase; PLEX=plasma exchange; PA=prior authorization



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