

Vyvgart (efgartigimod alfa-fcab) Medication Precertification Request

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(All fields must be completed and legible for precertification review.)

Aetna Precertification Notification

Phone: 1-866-752-7021 **FAX:** 1-888-267-3277

For Medicare Advantage Part B:

Phone: 1-866-503-0857 FAX: 1-844-268-7263

Precertification Requested By:		Phone:		Fax:	
A. PATIENT INFORMATION					
First Name:	Last Name:			DOB:	
Address:			City:		ZIP:
Home Phone: Work Phone:		Cell Phone:		Email:	
Patient Current Weight: lbs or kgs	Patient Height: inch	es orcms	Allergies:		
B. INSURANCE INFORMATION					
Aetna Member ID #:	Does patient have	other coverage?	☐ Yes ☐ No)	
Group #:	If yes, provide ID#:	-		:	
Insured:	Insured:				
Medicare: ☐ Yes ☐ No If yes, provide ID #:		Medicaid: Yes	☐ No If yes, p	orovide ID #:	
C. PRESCRIBER INFORMATION				_	
First Name:	Last Name:	•	(Check	One): M.D.	☐ D.O. ☐ N.P. ☐ P.A
Address:		City:		State:	ZIP:
Phone: Fax:	St Lic #:	NPI #:	DEA#	! :	UPIN:
Provider Email:	Office Contact Nan	ne:		Phone:	
Specialty (Check one): Neurologist Ot	her:				
D. DISPENSING PROVIDER/ADMINISTRATION	N INFORMATION				
Place of Administration:		Dispensing I	Provider/Pharr	nacy: Patient S	Selected choice
☐ Self-administered ☐ Physician's Office			☐ Physician's Office ☐ Retail Pharmacy		
Outpatient Infusion Center Phone:		=	☐ Specialty Pharmacy ☐ Other		•
Center Name:			-		
Agency Name:		Address:			
Administration code(s) (CPT):		Phone:		Fax:	
Address:		TIN:		PIN:	
E. PRODUCT INFORMATION					
Request is for: Vyvgart (efgartigimod alfa-fca	ıb) Dose:	Freque	ency:		
F. DIAGNOSIS INFORMATION - Please indica					
Primary ICD Code:				ner ICD Code:	
G. CLINICAL INFORMATION - Required clinical					
For Initiation Requests (clinical documentation			an precertifica	don requests.	
Generalized myasthenia gravis (gMG)					
☐ Yes ☐ No Is the requested drug being used	to treat a patient who is anti-ac	etylcholine receptor (A	AchR) antibody p	oositive?	
Please indicate the patient's Myasthenia Gravis Fo	oundation of America (MGFA) o	clinical classification:			
Please select: Class I Class II Class II					
Please indicate the patient's Myasthenia Gravis-Sp	•	, ,			
Yes No Is the MG-ADL score at least 50%				/	
Yes No Is the patient on a stable dose of a (at least 3 months of treatment) or		•			•
mycophenolate mofetil)?	nonsteroidar inimunosuppress	sive therapy (NSIST) (at least o month	s of treatment) (e.g., azatiliopilile,
For Continuation Requests (clinical documenta	tion required):				
☐ Yes ☐ No Is there evidence of unacceptable	toxicity or disease progression	while on the current	regimen?		
	sitive response to therapy (e.g.,	, improvement in MG-	ADL score, char	nges compared to	o baseline in
☐ Yes ☐ No Has the patient experienced a pos	MAO) 4-4-1 \0				
Quantitative Myasthenia Gravis (C	amG) total score)?				
	QMG) total score)?				
Quantitative Myasthenia Gravis (C	,			Dat	re:/
Quantitative Myasthenia Gravis (CH. ACKNOWLEDGEMENT	:uthorization of coverage of a	medical procedure	or service with	the intent to inj	jure, defraud or deceive