

Tegsedi® (inotersen) Medication **Precertification Request**

Page 1 of 1 (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 1-844-268-7263 FAX:

Please indicate:	ase indicate: Start of treatment: Start date// Continuation of therapy: Date of last treatment//								
		пегару. Басе	oriasi irealment	1 1					
	Requested By:			Phone:		Fa	ıx:		
A. PATIENT INFO	ORMATION								
First Name:			Last Name:	1		DOB:			
Address:				City:		State:	ZIP:		
Home Phone:		Work Phone:		Cell Phone:		Email:			
,	lbs orkg	s Height:	inches orcn	ns Allergies:					
B. INSURANCE	INFORMATION								
Aetna Member ID #:			Does patient have other coverage?						
			If yes, provide ID#: Carrier Name:						
Insured:			Insured:						
	s 🗌 No If yes, provid	le ID #:	Me	dicaid: Yes No	o If yes, prov	ide ID #:			
C. PRESCRIBER	RINFORMATION								
First Name:			Last Name:	1	Check One:		☐ D.O. ☐ N.P.	. ∐ P.A.	
Address:	T		1	City:		State:	ZIP:		
Phone:	Fax:		St Lic #:	NPI #:	DEA #:	T	UPIN:		
Provider Email:			Office Contact Name:			Phone:			
Specialty (Check	one): Neurologist	Other:							
D. DISPENSING	PROVIDER/ADMINIS	TRATION INFO	RMATION						
Place of Administration: Dispensing Provider/Pharmacy: (Patient selected choice)									
	red Physician's Off					ail Pharmacy			
•	sion Center Phone:			Specialty Pharmacy Other:					
Center Name: Home Infusion Center Phone:				Name:					
				Address:					
Agency Name:				Phone:		Fax:			
Address:				TIN:		PIN:			
E. PRODUCT IN	FORMATION								
Request is for Teg	gsedi (inotersen): Dose	e:		Frequency:					
	- :	•	ry ICD code and specif		licable.				
Primary ICD Code			,	Other ICD Code:					
-		d clinical inform	nation must be complete			a reguests			
			iation must be complete	ed in its <u>entirety</u> for all j	precertification	rrequests.			
	 (clinical documentated) of hereditary transthed 		d amyloidosis						
	=		=	TTR gene?					
Yes No Was the diagnosis confirmed by detection of a mutation in the TTR gene? Yes No Does the patient exhibit clinical manifestations of polyneuropathy of hereditary transthyretin-mediated amyloidosis (ATTR-FAP)									
(e.g., amyloid deposition in biopsy specimens, TTR protein variants in serum, progressive peripheral sensory-motor polyneuropathy)?									
	Yes No Is the patient a liver transplant recipient?								
	es No Will the requested medication be used in combination with patisiran (Onpattro), tafamidis (Vyndaqel, Vyndamax) or vutrisiran (Amvuttra)?								
Yes No Is the requested medication prescribed by or in consultation with any of the following: a) neurologist, b) geneticist, or c) physician specializing in the treatment of amyloidosis?								ian	
For Continuation		·							
r	neuropathy severity and	d rate of diseas orfolk Quality of	cial response to the req e progression as demor Life-Diabetic Neuropat	nstrated by the modifie	d Neuropathy	Impairmer	nt Scale+7 (mNIS	5+7)	
H. ACKNOWLED	DGEMENT								
	eted By (Signature Re	quired):				D	ate:/	1	
			tion of coverage of a m						

insurance act, which is a crime and subjects such person to criminal and civil penalties. The plan may request additional information or clarification, if needed, to evaluate requests.