

MEDICARE FORM Signifor LAR (pasireotide) **Medication Precertification Request**

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

For Ohio MMP: FAX: 1-855-734-9389 PHONE: 1-855-364-0974

For other lines of business: Please use other form

Note: Signifor LAR is nonpreferred for acromegaly. The preferred products are Sandostatin LAR and Somatuline

Please indicate:	Start of treatment:	Start date		

	—				Depot.		
	inuation of therapy, Date of						
recertification Requeste	d By:		Phone:	:	Fax:		
A. PATIENT INFORMATIO	N						
First Name:		Last Name:			DOB:		
Address:			City:		State:	ZIP:	
Home Phone:	Work Phone:		Cell Phone:		Email:		
Patient Current Weight:	lbs_orkgs_Patie	ent Height: inches	or <u>cms</u>	Allergies:	•		
B. INSURANCE INFORMA							Į
Aetna Member ID #:		Does patient have ot	ner coverage?	🗌 Yes 🗌 No			
Group #:		If yes, provide ID#:					
nsured:		Insured:					
Medicare: 🗌 Yes 🗌 No	If yes, provide ID #:	Me	edicaid: 🗌 Yes	No If yes, pro	vide ID #:		
C. PRESCRIBER INFORM	ATION						
First Name:		Last Name:		(Check C	0 <i>ne):</i> 🔲 M.D.	D.O. N.P. P.A	۱.
Address:			City:		State:	ZIP:	
Phone:	Fax:	St Lic #:	NPI #:	DEA #:		UPIN:	
Provider Email:	-	Office Contact Name			Phone:		
Specialty (Check one):	Endocrinologist 🗌 Othe	er:					
D. DISPENSING PROVIDE							ł
Place of Administration: Self-administered Outpatient Infusion Center	Physician's Office				cy: <i>Patient</i> Se ☐ Retail Pha ☐ Other		
Center Name:			_	-	_		
Home Infusion Center Agency Name:	Phone:		-				
Administration code(s) (CI	рт) .					710	
Address:						ZIP:	
City:	State:	ZIP:					
Phone:			-				
TIN:	PIN:		NPI:				
NPI:			-				
Request is for: 🗌 Signifor			Frequency:				
F. DIAGNOSIS INFORMAT	ION - Please indicate prima						
Primary ICD Code: 🗌		Secondary ICD Co			ICD Code:		
Please explain if there are an	ical documentation require referred for acromegaly. Th ent had prior therapy with Sign ent had a trial and failure, into statin LAR (octreotide acetate y other medical reason(s) that	d for all requests): e preferred products ar hifor LAR within the last 3 lerance, or contraindicatio) Somatuline Depot (e Sandostatin LAI 65 days? on to any of the follo (lanreotide)	R and Somatuline	Depot. at apply)	d for the patient's	
diagnosis? (select all that app Sandos	statin LAR (octreotide acetate	e) 🔲 Somatuline Depot ((lanreotide)				



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Patient First Name	Patient Last Name	Patient Phone	Patient DOB				
G CLINICAL INFORMATION (contin	(und) - Required clinical information mu	st he completed in its entirety for all pro	acertification requests				
G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its <u>entirety</u> for all precertification requests.							
☐ Acromegaly							
Please indicate the patient's pretreatment IGF-1 (insulin-like growth factor 1) level compared to the laboratory's reference normal range based on age							
and/or gender: 🔲 IGF-1 level is higher than the laboratory's normal range 🔛 IGF-1 level is lower than the laboratory's normal range							
☐ IGF-1 level falls within the laboratory's normal range							
Yes I No Has the patient had an inadequate or partial response to surgery?							
\longrightarrow Yes \square No Is there a clinical reason why the patient has not had surgery?							
Cushing's syndrome/disease							
☐ Yes ☐ No Did the patient have surgery that was not curative?							
\rightarrow Yes \square No Is the patient a candidate for surgery?							
For Continuation Requests (clinical documentation required for all requests):							
Acromegaly only:							
Please indicate how the patient's IGF-1 (insulin-like growth factor 1) level changed since initiation of therapy:							
□ IGF-1 level has increased □ IGF-1 level has decreased or normalized □ IGF-1 level has not changed							
H. ACKNOWLEDGEMENT							
Request Completed By (Signature	Required):		Date: / /				
Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent							

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.