SENTARA HEALTH PLANS

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-800-750-9692</u>. No additional phone calls will be necessary if all information (<u>including phone and fax #s</u>) on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process can be delayed.</u>

<u>Drug Requested</u> : select one drug below		
□ Egrifta SV®(tesamorelin)	□ Egrifta WR [™] (tesamorelin)	
MEMBER & PRESCRIBER INFOR	RMATION: Authorization may be delayed if incomplete.	
Member Name:		
Member Sentara #:	Date of Birth:	
Prescriber Name:		
Prescriber Signature:		
Office Contact Name:		
Phone Number: Fax Number:		
NPI #:		
DRUG INFORMATION: Authorizatio	n may be delayed if incomplete.	
Drug Name/Form/Strength:		
Dosing Schedule:	Length of Therapy:	
Diagnosis:	ICD Code, if applicable:	
Weight (if applicable):	Date weight obtained:	
 Quantity Limits: Egrifta SV Subcutaneous 2 mg single-p NDC 62064-241-30 	patient-use vials: 30 vials per 30 days	
• Egrifta WR Subcutaneous 11.6mg sings	le-patient-use vials: 4 vials per 28 days	
o NDC 62064-381-04		

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

Initial Authorization: 12 months

☐ Member is 18 years of age or older

(Continued on next page)

	Member has a diagnosis of Human Immunodeficiency Virus Infection with Lipodystrophy
	Medication is prescribed by or in consultation with an endocrinologist or a physician specializing in the treatment of HIV infection
	Member is currently receiving and adherent to antiretroviral therapy (verified by pharmacy paid claims)
	Prescribed therapy will <u>NOT</u> be used in combination with any form of growth hormone (somatropin) or IGF-1 (mecasermin)
	Provider will use tesamorelin to reduce excess abdominal visceral adipose tissue (VAT), and <u>NOT</u> for the following:
	Abdominal obesity in a patient without Human Immunodeficiency Virus (HIV) infection
	• Human Immunodeficiency Virus (HIV)-Related cachexia, weight loss, or fat distribution other than Lipodystrophy
	Member meets <u>ONE</u> of the following clinical indicators for abdominal lipodystrophy (submit documentation):
	☐ If female, waist circumference \geq 94 cm and waist-hip ratio \geq 0.88
	☐ If male, waist circumference \geq 95 cm and waist-hip ratio \geq 0.94
	Provider must submit documentation to confirm member has a body mass index (BMI) greater than 20 $\mbox{kg/m}^2$
	Member has no active malignancy (for example, a potential cancer which is being evaluated or a diagnosed cancer which is being treated)
	Member is NOT currently pregnant or breast-feeding
ıppo	uthorization: 12 months. Check below all that apply. All criteria must be met for approval. To ort each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be ded or request may be denied.
	Provider must submit documentation to confirm the member has exhibited a clear response in reduction of visceral adipose tissue measured by waist circumference or computed tomography (CT) scan
	Member does <u>NOT</u> demonstrate persistent elevated insulin-like growth factor 1 (IGF-1) levels (> 3 standard deviations above normal per the package insert)
	Member does <u>NOT</u> have unacceptable toxicity from the drug (e.g., severe injection site reactions, severe fluid retention, and severe hypersensitivity reactions)

$\label{eq:medication} \mbox{Medication being provided by Specialty Pharmacy-Proprium } \mbox{Rx}$

**Use of samples to initiate therapy does not meet step edit/ preauthorization criteria. **

*Previous therapies will be verified through pharmacy paid claims or submitted chart notes. *