

# AvMed

## PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST\*

**Directions:** The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **fax to 1-305-671-0200**. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If the information provided is not complete, correct, or legible, the authorization process can be delayed.**

**Drug Requested:** select one drug below

☐ **Egrifta SV<sup>®</sup>**(tesamorelin)

☐ **Egrifta WR<sup>™</sup>**(tesamorelin)

**MEMBER & PRESCRIBER INFORMATION:** Authorization may be delayed if incomplete.

**Member Name:** \_\_\_\_\_

**Member AvMed #:** \_\_\_\_\_ **Date of Birth:** \_\_\_\_\_

**Prescriber Name:** \_\_\_\_\_

**Prescriber Signature:** \_\_\_\_\_ **Date:** \_\_\_\_\_

**Office Contact Name:** \_\_\_\_\_

**Phone Number:** \_\_\_\_\_ **Fax Number:** \_\_\_\_\_

**NPI #:** \_\_\_\_\_

**DRUG INFORMATION:** Authorization 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-patient-use vials: 30 vials per 30 days
  - NDC 62064-241-30
- Egrifta WR Subcutaneous 11.6mg single-patient-use vials: 4 vials per 28 days
  - 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

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- ☐ 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 **NOT** 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 **NOT** 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 **ONE** 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 kg/m<sup>2</sup>
- ☐ 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

**Reauthorization: 12 months.** 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.

- ☐ 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 **NOT** demonstrate persistent elevated insulin-like growth factor 1 (IGF-1) levels ( $> 3$  standard deviations above normal per the package insert)
- ☐ Member does **NOT** have unacceptable toxicity from the drug (e.g., severe injection site reactions, severe fluid retention, and severe hypersensitivity reactions)

**Medication being provided by Specialty Pharmacy – Proprium 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.\****