

# SENTARA COMMUNITY PLAN (MEDICAID)

## 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 1-800-750-9692. 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:** Tryngolza<sup>™</sup> (olezarsen)

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

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

**Member Sentara #:** \_\_\_\_\_ **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 Limit:** 80 mg/0.8 mL autoinjector – one autoinjector per 30 days

**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
- ☐ Prescribed by or in consultation with a cardiologist, endocrinologist, or a specialist experienced in treating severe hypertriglyceridemia
- ☐ Member has a diagnosis of Familial Chylomicronemia Syndrome (FCS) that is supported by genetic testing showing biallelic pathogenic variants in FCS-causing genes (LPL, LMF1, GPIHBP1, APOC2, APOA5) (**submit results of genetic testing**)
- ☐ Member has fasting triglyceride level  $\geq 880$  mg/dL (**submit lab results from the past 30 days**)
- ☐ Requested medication will be used as an adjunct to a low-fat diet ( $\leq 20$  g of fat per day)

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**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.

- ☐ Member has experienced positive clinical response from the medication as demonstrated by improvement in fasting triglyceride levels (**submit lab results from the past 90 days**)
- ☐ Requested medication will continue to be used as an adjunct to a low-fat diet ( $\leq 20$  g of fat per day)
- ☐ Member has **NOT** experienced serious adverse events related to the medication (thrombocytopenia, hypersensitivity to olezarsen)

**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.\****