

# 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:** Wegovy® (semaglutide)- GLP-1 Receptor Agonists  
for Metabolic Dysfunction-Associated Steatohepatitis (MASH)

**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: \_\_\_\_\_

**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.

- **FDA indicated medications only**
- **Must be prescribed by or in consultation with a hepatologist or gastroenterologist or other provider specializing in liver disease for the member to receive authorization.**

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**Initial Request Requirements: 6 months**

- ☐ The member is 18 years of age or older; **AND**
- ☐ The medication is prescribed by or in consultation with a hepatologist, gastroenterologist or other provider specializing in liver disease; **AND**
- ☐ The member has a diagnosis of MASH with results of baseline liver biopsy or noninvasive tests demonstrating the presence of stage F2 or F3 fibrosis by at least **ONE** of the following:
  - ☐ Liver biopsy; **OR**
  - ☐ Noninvasive tests (such as transient elastography, Fibroscan, or magnetic resonance elastography) performed within the last 6 months; **AND**
- ☐ The member has a BMI  $\geq 18.5$  kg/m<sup>2</sup>; **AND**
- ☐ The provider attests that the member received individualized healthy lifestyle counseling; **AND**
- ☐ The member does not have an A1C of  $>9.5\%$  **AND**
- ☐ The member does not have known or suspected excessive consumption of alcohol according to the CDC's guidance; **AND**
- ☐ The member does not have hepatic decompensation or a MELD score of  $> 12$  at screening; **AND**
- ☐ The member does not have pancreatitis, acute suicidal behavior/ideation, personal or family history of medullary thyroid cancer or multiple endocrine neoplasia 2 syndrome; **AND**
- ☐ The member is not concurrently on another GLP-1 receptor agonist

**Renewal requests: 12 months**

- ☐ The member has experienced clinical improvement on the requested medication
- ☐ The member is being treated with a maintenance dosage of the requested drug

***\*\*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.\****