

SENTARA HEALTH PLAN

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: Myalept® (metreleptin)

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: _____

DEA OR NPI #: _____

DRUG INFORMATION: Complete **all** information below or authorization may be delayed.

Drug Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

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.

INITIATION AND CONTINUATION OF TREATMENT – All boxes below must be checked to qualify.

- ☐ Member has a leptin deficiency as defined as **(a copy of fasting laboratory leptin assay results is required for approval):**
 - ☐ <4.0 ng/mL fasting leptin for females
 - ☐ <3.0 ng/mL fasting leptin for males
- ☐ Member has a diagnosis of **(choose indication):**
 - ☐ Acquired generalized lipodystrophy
 - ☐ Congenital generalized lipodystrophy
- ☐ Member has a concurrent condition of:

(Continued on next page)

- ☐ Diabetes mellitus or insulin resistance and failed 30-day trial of (submit chart notes):
☐ Metformin, total daily dose of: _____

AND

- ☐ High-dose insulin or insulin pump
☐ Hypertriglyceridemia and failed 30-day trial of (submit chart notes):
☐ Low-fat diet and/or dietary restrictions

AND

- ☐ Fenofibrate or fenofibrate derivative

OR

- ☐ Niacin or omega-3 fatty acid

OR

- ☐ Atorvastatin, simvastatin, pravastatin, rosuvastatin

OR

- ☐ Other therapy of (please specify): _____

INITIATION OF TREATMENT (submit all labs)	<u>REAUTHORIZATION</u> (submit all labs)
HbA1c%	HbA1c%
Fasting glucose _____ mg/dL	Fasting glucose _____ mg/dL
Triglyceride _____ mg/dL	Triglyceride _____ mg/dL
Patient weight _____ kg	Patient weight _____ kg
	Has member experienced clinical improvement or metabolic stabilization while using this medication? (submit chart notes to verify response) ☐ Yes ☐ No

If approved, response to initial treatment will be **assessed after 4 months**, then **quarterly reassessment** will be required for continued approval.

Medication being provided by Specialty Pharmacy - PropriumRx

****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.