

# HEALTHY U MEDICAID

## PRIOR AUTHORIZATION REQUEST FORM SANDOSTATIN LAR®

**For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U Prior Authorization Department.**

- For **Medical Pharmacy** please fax requests to: 801-213-1547
- For **Retail Pharmacy** please fax requests to: 385-425-4052

**Failure to submit clinical documentation to support this request will result in a dismissal of the request.**

If you have prior authorization questions, please call for Pharmacy Customer Service for assistance at 385-425-5094

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

|                |              |                 |
|----------------|--------------|-----------------|
| Date:          | Member Name: | ID#:            |
| DOB:           | Gender:      | Physician:      |
| Office Phone:  | Office Fax:  | Office Contact: |
| Height/Weight: | HCPCS Code:  |                 |

**Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.**

**Product being requested:**  Sandostatin® LAR (octreotide)

Dosing/Frequency: \_\_\_\_\_

### If the request is for reauthorization, proceed to reauthorization section

| Questions  | Yes                      | No                       | Comments/Notes                      |
|--|--------------------------|--------------------------|-------------------------------------|
| 1. Has the member had a clinical response and tolerance to immediate-release octreotide prior to depot injection use?      | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| <b>ACROMEGALY</b>  |                          |                          |                                     |
| 2. Has the member had an inadequate response or contraindication to surgery or radiation?                                  | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| 3. Has the member had an inadequate response or contraindication to a dopamine agonist (i.e., bromocriptine, cabergoline)? | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| <b>METASTATIC CARCINOID TUMORS</b>   |                          |                          |                                     |
| 1. Does the member have severe diarrhea and flushing associated with metastatic carcinoid tumors?                          | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| <b>VASOACTIVE INTESTINAL PEPTIDE TUMOR (VIPoma)</b>  |                          |                          |                                     |
| 1. Does the member have profuse watery diarrhea associated with a Vasoactive Intestinal Peptide Tumor (VIPoma)?            | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| <b>Gastrointestinal Arterio-Venous Malformations (HEYDE'S SYNDROME)</b>  |                          |                          |                                     |
| 1. Is the request for gastrointestinal arteriovenous malformations (e.g. Heyde's Syndrome)?                                | <input type="checkbox"/> | <input type="checkbox"/> |                                     |

| NEUROENDOCRINE TUMORS  |                          |                          |                                     |
|--|--------------------------|--------------------------|-------------------------------------|
| 1. Is the request for neuroendocrine tumors and in accordance with NCCN guidelines?  | <input type="checkbox"/> | <input type="checkbox"/> |                                     |
| REFRACTORY DIARRHEA ASSOCIATED WITH ACUTE GRAFT VERSUS HOST DISEASE OR CHEMOTHERAPY  |                          |                          |                                     |
| 1. Is the request for refractory diarrhea associated with acute graft versus host disease or chemotherapy?   | <input type="checkbox"/> | <input type="checkbox"/> |                                     |
| HIGH OUTPUT FISTULAS   |                          |                          |                                     |
| 1. Is the request for high output fistulas?  | <input type="checkbox"/> | <input type="checkbox"/> |                                     |
| REAUTHORIZATION  |                          |                          |                                     |
| 1. Is the request for reauthorization of therapy?  | <input type="checkbox"/> | <input type="checkbox"/> |                                     |
| 2. Has the therapy shown to be effective with a clinically significant response to therapy?  | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| 3. Does the member show a continued medical need for the therapy?  | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| <b>What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.</b> |                          |                          |                                     |
| Additional information:  |                          |                          |                                     |
| Physician's Signature:   |                          |                          |                                     |

**\*\*Failure to submit clinical documentation to support this request will result in a dismissal of the request.\*\***

Policy PHARM-HU-066  
 Origination Date: 01/01/2022  
 Reviewed/Revised Date: 09/13/2023  
 Next Review Date: 09/13/2024  
 Current Effective Date: 10/01/2023

**Confidentiality Notice**

This document and any accompanying document contain confidential information and is intended for the use of the individual or entity named on this transmission sheet. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution or the taking of any action in reliance on the contents of this information is strictly prohibited and the document should be returned to this office immediately. If you have received this facsimile in error, please notify us by telephone immediately and destroy document received.