



Lutathera

Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720.** If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____
Specialty: _____ **NPI#:** _____
Physician Office Telephone: _____ **Physician Office Fax:** _____

Referring Provider Info: Same as Requesting Provider
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Rendering Provider Info: Same as Referring Provider Same as Requesting Provider
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

Patient Weight: _____ kg
Patient Height: _____ cm

Please indicate the place of service for the requested drug:

- Ambulatory Surgical Home Off Campus Outpatient Hospital
 On Campus Outpatient Hospital Office Pharmacy

Criteria Questions:

1. What is the diagnosis?
 Neuroendocrine tumors of the gastrointestinal (GI) tract (carcinoid tumors)
 Neuroendocrine tumors of the pancreas
 Neuroendocrine tumors of the lung and thymus (carcinoid tumors)
 Poorly controlled carcinoid syndrome
 Pheochromocytoma/paraganglioma
 Other
2. What is the ICD-10 code? _____
3. Will the patient receive more than 4 doses total of the requested drug? Yes No

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. Lutathera SGM – 06/2021.

**CVS Caremark Specialty Pharmacy • 2211 Sanders Road NBT-6 • Northbrook, IL 60062
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4. Does the patient have any of the following? **Indicate ALL that apply.**
- | | |
|--|--|
| <input type="checkbox"/> Clinically significant tumor burden | <input type="checkbox"/> Low grade (typical) histology |
| <input type="checkbox"/> Distant metastases | <input type="checkbox"/> Distant metastatic disease |
| <input type="checkbox"/> Evidence of progression | <input type="checkbox"/> Intermediate grade (atypical) histology |
| <input type="checkbox"/> Locoregional unresectable disease | <input type="checkbox"/> Progressive locoregional advanced disease |
| <input type="checkbox"/> Symptomatic disease | <input type="checkbox"/> None of the above |

Complete the following section based on the patient's diagnosis, if applicable.

Section A: NETs of GI, Pancreas or Lung/Thymus

5. Are the patient's tumors somatostatin receptor-positive? **ACTION REQUIRED: If 'Yes', attach documentation supporting positive somatostatin receptor status as detected by somatostatin receptor-based imaging.** Yes No/unknown
6. Has the patient experienced disease progression on octreotide (Sandostatin, Sandostatin LAR) or lanreotide (Somatuline Depot)? Yes No

Section B: Carcinoid Syndrome, Poorly Controlled

7. Does the patient have somatostatin receptor-positive neuroendocrine tumors of the gastrointestinal tract, lung or thymus? **ACTION REQUIRED: If 'Yes', attach documentation supporting positive somatostatin receptor status as detected by somatostatin receptor-based imaging.** Yes No
8. Has the patient experienced progression on octreotide (Sandostatin, Sandostatin LAR) or lanreotide (Somatuline Depot)? Yes No
9. How will Lutathera be used?
- | |
|---|
| <input type="checkbox"/> In combination with octreotide LAR (Sandostatin LAR) |
| <input type="checkbox"/> In combination with lanreotide (Somatuline Depot) |
| <input type="checkbox"/> In combination with telotristat (Xermelo), <i>skip to 11</i> |
| <input type="checkbox"/> None of the above |
10. Does the patient have persistent symptoms (i.e., flushing, diarrhea)? Yes No *Nofurther questions*
11. Does the patient have persistent diarrhea? Yes No

Section C: Pheochromocytoma/Paraganglioma

12. Does the patient have somatostatin receptor-positive pheochromocytoma/paraganglioma? **ACTION REQUIRED: If 'Yes', attach documentation supporting positive somatostatin receptor status as detected by somatostatin receptor-based imaging.** Yes No/unknown

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____

Prescriber or Authorized Signature

Date (mm/dd/yy)

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