

**Sandostatin Injection /Sandostatin LAR Depot (for Maryland only)**  
**Prior Authorization Request**

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

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 **1-866-814-5506**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect<sup>®</sup> 1-800-237-2767.

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

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

**Additional Demographic Information:**

*Patient Weight:* \_\_\_\_\_ *kg*  
*Patient Height:* \_\_\_\_\_ *ft* \_\_\_\_\_ *inches*

**Criteria Questions:**

1. Which drug is being prescribed?
 

<input type="checkbox"/> Sandostatin Injection	<input type="checkbox"/> Sandostatin LAR Depot
<input type="checkbox"/> Octreotide acetate injection (generic)	<input type="checkbox"/> Other _____
  
2. What is the patient's diagnosis?
  - Acromegaly
  - Meningioma
  - Thymomas and thymic carcinoma
  - Neuroendocrine tumors of the gastrointestinal tract (carcinoid tumors)
  - Neuroendocrine tumors of the thymus (carcinoid tumors)
  - Neuroendocrine tumors of the lung (carcinoid tumors)
  - Pancreatic neuroendocrine tumors
  - Adrenal gland neuroendocrine tumors
  - Poorly differentiated (high-grade) neuroendocrine tumors/Large or small cell tumors (excluding lung)
  - Congenital hyperinsulinism/persistent hyperinsulinemic hypoglycemia of infancy (CHI/PHHI)

*Document patient's age:* \_\_\_\_\_ month(s) or year(s)

- Other \_\_\_\_\_
  
3. What is the ICD-10 code? \_\_\_\_\_
  
4. Would the prescriber like to request an override of the step therapy requirement?  Yes  No *If No, skip to diagnosis section.*
  
5. Has the member received the medication through a pharmacy or medical benefit within the past 180 days?
 

<input type="checkbox"/> Yes	<input type="checkbox"/> No
------------------------------	-----------------------------

**ACTION REQUIRED: *Please provide documentation to substantiate the member had a prescription paid for within the past 180 days (i.e. PBM medication history, pharmacy receipt, EOB etc.)***

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. Sandostatin CareFirst – 3/2016.

CVS Caremark is an independent company that provides pharmacy benefit management services to CareFirst and BlueChoice members.

CareFirst BlueCross BlueShield is the shared business name of CareFirst of Maryland, Inc. and Group Hospitalization and Medical Services, Inc. CareFirst BlueCross BlueShield and CareFirst BlueChoice, Inc. are both independent licensees of the Blue Cross and Blue Shield Association. The Blue Cross and Blue Shield Names and Symbols are registered trademarks of the Blue Cross and Blue Shield Association. © Registered trademark of CareFirst of Maryland, Inc.

6. Is the medication effective in treating the member's condition?  Yes  No *Continue to diagnosis section and complete this form in its entirety.*

**Complete the following section based on the patient's diagnosis.**

Section A: Acromegaly

7. Does the patient have clinical evidence of acromegaly (e.g., frontal bossing, coarse facial features, thick lips, protruding jaw with widely spaced teeth, large hands and feet)?  Yes  No
8. Is the patient currently on the requested medication?  Yes  No *If No, skip to #11*
9. What is the **current** IGF-1 level? \_\_\_\_\_ **ACTION REQUIRED: Attach lab documentation of current IGF-1 level.**
10. How has the patient's IGF-1 level changed since initiation of therapy? *Indicate below and no further questions.*  
 Increased  Decreased or normalized  No change
11. What is the **pretreatment** IGF-1 level? \_\_\_\_\_ **ACTION REQUIRED: Attach lab documentation of pretreatment IGF-1 level.**
12. How does the patient's IGF-1 level compare to the laboratory's reference normal range based on age and/or gender? (Note: The normal range varies based on the laboratory performing the analysis. One must obtain lab-specific values to make this determination.)  
 IGF-1 level is **higher** than the laboratory's normal range  
 IGF-1 level is **lower** than the laboratory's normal range  
 IGF-1 level **falls within** the laboratory's normal range
13. Has the patient had an inadequate or partial response to surgery or radiotherapy?  
*If Yes, no further questions*  Yes  No
14. Is there a clinical reason why the patient has not had surgery or radiotherapy?  Yes  No
15. What is the clinical reason for not having surgery or radiotherapy?  
 Patient is medically unstable (poor surgical candidate)  
 Patient is at high risk for complications of anesthesia because of airway difficulties  
 Patient has major systemic manifestations of acromegaly including cardiomyopathy, severe hypertension and uncontrolled diabetes  
 Patient refuses surgery or prefers the medical option over surgery  
 Lack of an available skilled surgeon  
 Other \_\_\_\_\_

Section B: Meningioma

16. Is the disease recurrent or progressive?  Yes  No
17. Is the disease unresectable?  Yes  No
18. Is the disease refractory to radiation therapy?  Yes  No
19. What is the tumor somatostatin receptor status?  Positive  Negative  Unknown

Section C: Thymomas and Thymic Carcinoma

20. Does the patient have unresectable disease? *If Yes, skip to #22*  Yes  No
21. Was there residual disease following resection?  Yes  No
22. Does the patient have locally advanced, advanced, or recurrent disease?  Yes  No
23. Has the patient progressed on at least one prior chemotherapy regimen?  Yes  No
24. Does the patient experience symptoms of carcinoid syndrome (eg, skin flushing, diarrhea)?  
*If Yes, no further questions*  Yes  No
25. What is the tumor somatostatin receptor status?  Positive  Negative  Unknown

Section D: Neuroendocrine Tumors of the Gastrointestinal Tract (Carcinoid Tumors)

26. Does the patient have distant metastases? *If Yes, no further questions*  Yes  No

27. Is the disease unresectable? *If Yes, no further questions*  Yes  No
28. What is the primary site of the tumor?  
 Gastric  Jejunal/ileal/colon  Duodenal  Appendix  Rectal  Other \_\_\_\_\_
29. What is the tumor size? \_\_\_\_\_ centimeters
30. Does the patient have hypersecretion of gastrin (eg, Zollinger-Ellison syndrome)?  Yes  No

Section E: Neuroendocrine Tumors of the Thymus (Carcinoid Tumors)

31. Does the patient have distant metastases? *If Yes, no further questions*  Yes  No
32. Is the disease unresectable?  Yes  No

Section F: Neuroendocrine Tumors of the Lung (Carcinoid Tumors)

33. Does the patient have distant metastases? *If Yes, no further questions*  Yes  No
34. What is the tumor grade?  
 High-grade neuroendocrine carcinoma (eg, large cell neuroendocrine carcinoma [LCNEC], small cell carcinoma, combined SCLC and NSCLC)  
 Intermediate-grade neuroendocrine carcinoma (atypical carcinoid)  
 Low-grade neuroendocrine carcinoma (typical carcinoid)  
 Other \_\_\_\_\_
35. What is the disease stage?  
 I (IA, IB)  II (IIA, IIB)  III (IIIA, IIIB)  IV, *skip to #38*
36. Is the disease Stage IIIB?  Yes  No
37. Is the tumor stage T4 due to multiple lung nodules?  Yes  No
38. Does the patient experience symptoms of carcinoid syndrome (eg, flushing, diarrhea)?  
*If Yes, no further questions*  Yes  No
39. What is the tumor somatostatin receptor status?  Positive  Negative  Unknown

Section G: Pancreatic Neuroendocrine Tumors

40. What is the tumor type?  
 Gastrinoma, *skip to #43*  
 Glucagonoma, *skip to #43*  
 Vasoactive intestinal peptide tumor (VIPoma), *skip to #43*  
 Insulinoma  
 Non-functioning pancreatic tumors  
 Somatostatinoma  
 Pancreatic polypeptidoma (PPoma)  
 Cholecystokininoma (CCKoma)  
 ACTH-secreting pancreatic neuroendocrine tumor  
 Parathyroid hormone-related protein (PTHrp)-secreting pancreatic neuroendocrine tumor  
 Other \_\_\_\_\_
41. Does the patient have distant metastases? *If Yes, skip to #43*  Yes  No
42. Is the disease unresectable?  Yes  No
43. Does the patient experience hormone-related symptoms (eg, fasting or nocturnal hypoglycemia for insulinomas, recurrent peptic ulcers for gastrinomas, flushing, diarrhea)? *If Yes, no further questions*  Yes  No
44. What is the tumor somatostatin receptor status?  Positive  Negative  Unknown

Section H: Adrenal Gland Neuroendocrine Tumors

45. Does the patient have a diagnosis of non-adrenocorticotrophic hormone (non-ACTH) dependent Cushing's syndrome?  Yes  No
46. What is the tumor somatostatin receptor status?  Positive  Negative  Unknown
47. Is the cortisol production symmetric?  Yes  No

48. What is the tumor size? \_\_\_\_\_ centimeters

Section I: Poorly Differentiated (high-grade) Neuroendocrine Tumors/Large or Small Cell Tumors (excluding lung)

49. Does the patient have metastatic disease? *If Yes, skip to #51*  Yes  No

50. Is the disease unresectable?  Yes  No

51. What is the tumor somatostatin receptor status?  Positive  Negative  Unknown

52. Does the patient experience hormone-related symptoms?  Yes  No

*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)**