Standard Opt-in	PDPD	Marketplace	Medical Benefit	1	Medicare Part B
Standard Opt-out	ACSF	ммт	Medical Benefit: Biosimilars First		Medicare Part B: Biosimilars First
VF	Balanced	Medical Benefit: Managed Medicaid	Medical Benefit: Add-on		Medicare Part B: Add-on

Reference #
4255-D

# **POLICY Document for SANDOSTATIN LAR**

The overall objective of this policy is to support the appropriate and cost effective use of the medication, specific to use of preferred medication options, lower cost site of care and overall clinically appropriate use. This document provides specific information to each section of the overall policy.

#### Section 1: Preferred Product

• Policy information specific to preferred medications

#### Section 2: Clinical Criteria

• Policy information specific to the clinical appropriateness for the medication

# **Section 1: Preferred Product**

# EXCEPTIONS CRITERIA ACROMEGALY PRODUCTS

#### PREFERRED PRODUCTS: SOMATULINE DEPOT, SANDOSTATIN LAR

#### **POLICY**

This policy informs prescribers of preferred products and provides an exception process for targeted products through prior authorization.

# I. PLAN DESIGN SUMMARY

This program applies to the acromegaly products specified in this policy. Coverage for a targeted product is provided based on clinical circumstances that would exclude the use of the preferred products and may be based on previous use of a product. The coverage review process will ascertain situations where a clinical exception can be made. This program applies to members who are new to treatment with the targeted product for the first time.

Each referral is reviewed based on all utilization management (UM) programs implemented for the client.

#### **Table. Acromegaly Products**

	Product(s)
Preferred	Somatuline Depot (lanreotide)
	Sandostatin LAR (octreotide acetate for injectable suspension)
Targeted	Signifor LAR (pasireotide)
	Somavert (pegvisomant)

# II. EXCEPTION CRITERIA

This program applies to members requesting treatment for an indication that is FDA-approved for both of the preferred products.

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	ndard t-in PDPD	Marketplace	Medical Benefit	1	Medicare Part B
	ndard t-out ACSF	ммт	Medical Benefit: Biosimilars First		Medicare Part B: Biosimilars First
VF	Balanced	Medical Benefit: Managed Medicaid	Medical Benefit: Add-on		Medicare Part B: Add-on

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Coverage for a targeted product is provided when either of the following criteria is met:

- A. Member has received treatment with the requested targeted product in the past 365 days.
- B. Member has a documented inadequate response or intolerable adverse event to any of the preferred products.

#### REFERENCES

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# **Section 2: Clinical Criteria**

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# STANDARD MEDICARE PART B MANAGEMENT

SANDOSTATIN LAR (octreotide acetate for injectable suspension)

# **POLICY**

# I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

#### A. FDA-Approved Indications

Sandostatin LAR Depot is indicated in patients in whom initial treatment with Sandostatin injection has been shown to be effective and tolerated.

 Indicated for long-term maintenance therapy in acromegalic patients who have had an inadequate response to surgery and/or radiotherapy, or for whom surgery and/or radiotherapy is not an option.

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Standard Opt-in	PDPD	Marketplace	Medical Benefit	1	Medicare Part B
Standard Opt-out	ACSF	ммт	Medical Benefit: Biosimilars First		Medicare Part B: Biosimilars First
VF	Balanced	Medical Benefit: Managed Medicaid	Medical Benefit: Add-on		Medicare Part B: Add-on

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- Indicated for long-term treatment of the severe diarrhea and flushing episodes associated with metastatic carcinoid tumors.
- Indicated for long-term treatment of the profuse watery diarrhea associated with vasoactive intestinal peptide (VIP)-secreting tumors.

#### B. Compendial Uses

- 1. Neuroendocrine tumors (NETs)
  - NETs of the gastrointestinal (GI) tract, lung, and thymus (carcinoid tumors) or unresected primary gastrinoma
  - o. NETs of the pancreas
- 2. Pheochromocytoma/paraganglioma
- 3. Meningiomas
- 4. Thymomas and thymic carcinomas
- 5. Treatment of diarrhea in acquired immunodeficiency syndrome (AIDS)
- 6. Bowel obstruction due to peritoneal carcinomatosis
- 7. Postgastrectomy dumping syndrome
- 8. Pediatric hypothalamic obesity
- 9. Hepatocellular carcinoma
- 10. Pancreatic pleural effusion
- 11. Pituitary adenoma
- 12. Graves' ophthalmopathy
- 13. von Willebrand disorder

All other indications will be assessed on an individual basis. Submissions for indications other than those enumerated in this policy should be accompanied by supporting evidence from Medicare approved compendia.

### II. CRITERIA FOR INITIAL APPROVAL

## A. Acromegaly

Authorization of 24 months may be granted for the treatment of acromegaly when all of the following criteria are met:

- 1. Member has a high pretreatment insulin-like growth factor-1 (IGF-1) level for age and/or gender based on the laboratory reference range.
- 2. Member had an inadequate or partial response to surgery or radiotherapy OR there is a clinical reason why the member has not had surgery or radiotherapy.

# B. Carcinoid syndrome

Authorization of 12 months may be granted for treatment of carcinoid syndrome.

#### C. Vasoactive intestinal peptide tumors (VIPomas)

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Standard Opt-in	PDPD	Marketplace	Medical Benefit	1	Medicare Part B
Standard Opt-out	ACSF	ммт	Medical Benefit: Biosimilars First		Medicare Part B: Biosimilars First
VF	Balanced	Medical Benefit: Managed Medicaid	Medical Benefit: Add-on		Medicare Part B: Add-on

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Authorization of 12 months may be granted for management of symptoms related to hormone hypersecretion of VIPomas.

#### D. Neuroendocrine tumors (NETs)

- Authorization of 12 months may be granted for treatment of NETs of the gastrointestinal (GI) tract, lung, thymus (carcinoid tumors) or unresected primary gastrinoma.
- 2. Authorization of 12 months may be granted for treatment of NETs of the pancreas.

### E. Pheochromocytoma and paraganglioma

Authorization of 12 months may be granted for treatment of pheochromocytoma/paraganglioma.

# F. Meningiomas

Authorization of 12 months may be granted to members for treatment of unresectable meningioma.

#### G. Thymomas and thymic carcinomas

Authorization of 12 months may be granted for treatment of thymoma and thymic carcinoma.

#### H. Bowel obstruction due to peritoneal carcinomatosis

Authorization of 12 months may be granted for treatment of bowel obstruction due to peritoneal carcinomatosis.

# l. Postgastrectomy dumping syndrome

Authorization of 12 months may be granted for treatment of postgastrectomy dumping syndrome.

#### J. Hepatocellular carcinoma

Authorization of 12 months may be granted for treatment of hepatocellular carcinoma.

# K. Pituitary adenoma

Authorization of 12 months may be granted for treatment of pituitary adenoma.

#### L. Diarrhea in acquired immunodeficiency syndrome (AIDS)

Authorization of 12 months may be granted for treatment of diarrhea in a member with acquired immunodeficiency syndrome (AIDS).

# M. Pediatric hypothalamic obesity

Authorization of 12 months may be granted for treatment of hypothalamic obesity in a pediatric member.

# N. Pancreatic pleural effusion

Authorization of 12 months may be granted for treatment of pancreatic pleural effusion.

# O. Graves' ophthalmopathy

Authorization of 12 months may be granted for treatment of Graves' ophthalmopathy.

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Opt-out	ACSF	MMT	Biosimilars First		Biosimilars First
		Medical Benefit:	Medical Benefit:		Medicare Part B:
VF	Balanced	Managed Medicaid	Add-on		Add-on

Reference #	
4255-D	

#### P. von Willebrand disorder

Authorization of 12 months may be granted for treatment of von Willebrand disorder.

#### **III. CONTINUATION OF THERAPY**

All members (including new members) requesting authorization for continuation of therapy must be currently receiving therapy with the requested agent.

Authorization for 12 months may be granted when all of the following criteria are met:

- 1. The member is currently receiving therapy with Sandostatin LAR.
- Sandostatin LAR is being used to treat an indication enumerated in Section II.
- 3. The member is receiving benefit from therapy.

# **REFERENCES:**

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- Somatuline Depot [package insert]. Basking Ridge, NJ: Ipsen Biopharmaceuticals, Inc.; June 2019.
- Sandostatin LAR Depot [package insert]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; April 2019.
- 3. Signifor LAR [package insert]. East Hanover, NJ: Novartis Pharmaceuticals Company; April 2019.
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Standard Opt-out	ACSF	MMT	Medical Benefit: Biosimilars First		Medicare Part B: Biosimilars First
VF	Balanced	Medical Benefit: Managed Medicaid	Medical Benefit: Add-on		Medicare Part B: Add-on

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