

## octreotide (Sandostatin IV or LAR Depot)

### Commercial Medical Benefit Drug Policy

For oncology-related indications, medical necessity criteria can be found here: [Blue Shield Oncology-Related Medication Policies](#).

For PPO, Direct Contract HMO, and when applicable, ASO, and Shared Advantage: Please access Evolent's [CarePro Provider Portal](#) to submit your request.

#### Place of Service

##### *Sandostatin IV*

Home Infusion Administration

Hospital Administration

Infusion Center Administration

Outpatient Facility Infusion Administration

##### *Sandostatin LAR Depot - IM only*

Home Infusion Administration

Hospital Administration

Infusion Center Administration

Office Administration

Outpatient Facility Infusion Administration

### Drug Details

**USP Category:** HORMONAL AGENTS, SUPPRESSANT (ADRENAL OR PITUITARY)

**Mechanism of Action:** Octreotide is a synthetic polypeptide structurally and pharmacologically related to somatostatin (growth hormone [somatropin] release inhibiting factor)

#### **HCPCS:**

J2353:Injection, octreotide, depot form for intramuscular injection, 1 mg

J2354:Injection, octreotide, non-depot form for subcutaneous or intravenous injection, 25 mcg

#### **How Supplied:**

- Sandostatin:
  - 50 mcg, 100 mcg, or 500 mcg single-dose 1 mL ampuls
  - 1000 mcg or 5000 mcg multi-dose vial
- Sandostatin LAR: 10 mg, 20 mg or 30 mg single-use vials

### Condition(s) listed in policy (see coverage criteria for details)

- Acromegaly
- AIDS-Associated Diarrhea
- Bleeding Esophageal Varices
- Chemotherapy-Induced Diarrhea, treatment
- Cryptosporidiosis
- Dumping Syndrome
- Lymphorrhagia
- Malignant Intestinal Obstruction
- Pancreatitis, necrotizing

- Pituitary Adenomas (TSH-Secreting)
- Polycystic Ovary Syndrome (PCOS)
- Prevention of Postoperative Complications of Pancreatic Surgery
- Radiation-Induced Diarrhea
- Zollinger-Ellison Syndrome (Gastrinoma)

Any condition not listed in this policy requires a review to confirm it is medically necessary. For conditions that have not been approved for intended use by the Food and Drug Administration (i.e., off-label use), the criteria outlined in the Health and Safety Code section 1367.21 must be met.

### **Special Instructions and Pertinent Information**

Provider must submit documentation (such as office chart notes, lab results or other clinical information) to ensure the member has met all medical necessity requirements.

The member's specific benefit may impact drug coverage. Other utilization management processes, and/or legal restrictions may take precedence over the application of this clinical criteria.

For billing purposes, drugs must be submitted with the drug's assigned HCPCS code (as listed in the drug policy) and the corresponding NDC (national drug code). An unlisted, unspecified, or miscellaneous code should not be used if there is a specific code assigned to the drug.

Sandostatin given by subcutaneous injection: Refer to the "Self-Administered Drugs" policy

Sandostatin given intravenously and Sandostatin LAR Depot are managed under the Medical Benefit.

### **Coverage Criteria**

**The following condition(s) require Prior Authorization/Preservice.**

#### **Acromegaly**

**Meets medical necessity if all the following are met:**

1. Being prescribed by or recommended by an endocrinologist

#### **Covered Doses:**

Sandostatin IV: Up to 1500 mcg per day

Sandostatin LAR Depot IM: Up to 40 mg every 4 weeks

#### **Coverage Period:**

Sandostatin IV:

Initial: 2 weeks then reassess response

Reauthorization: Every 6 months depending on patient response

Sandostatin LAR Depot IM: indefinitely

#### **ICD-10:**

E22.0, E34.4

#### **AIDS-Associated Diarrhea**

**Meets medical necessity if all the following are met:**

octreotide (Sandostatin IV or LAR Depot)

1. Patient is currently stable on an anti-retroviral therapy (ART) regimen for  $\geq 1$  month
2. Provider attestation for infectious cause for diarrhea symptoms or other treatable causes (e.g. malabsorption, underlying GI disease requiring treatment) have been ruled out
3. Patient has inadequate response, intolerance, or contraindication to regular use of generic Lomotil or loperamide (OTC)
4. Patient has inadequate response, intolerance, or contraindication to Mytesi (crofelemer)

**Covered Doses:**

Sandostatin IV: Up to 1500 mcg/day

Sandostatin LAR IM: Up to 40 mg every 4 weeks

**Coverage Period:**

Yearly dependent upon patient response

**ICD-10:**

K52.2, K52.89, R19.7 + HIV infection B20 or B97.35

**Prevention of Postoperative Complications of Pancreatic Surgery**

**Meets medical necessity if all the following are met:**

1. Being used to prevent complications of pancreatic surgery (i.e., abscess formation, sepsis, acute pancreatitis, pancreatic fistula, and peripancreatic fluid collection)

**Covered Doses:**

Sandostatin IV: Up to 300 mcg per day

**Coverage Period:**

Sandostatin IV:

Initial: 2 weeks then reassess response

Reauthorization: Every 6 months depending on patient response

**ICD-10:**

BF1, ODB, ODJ, OWJ, OF1, OF5, OF7, OF8, OF9, OFB, OFC, OFF, OFH, OFJ, OFL, OFM, OFP-OFW

**The following condition(s) DO NOT require Prior Authorization/Preservice if ALL its parameters are met, otherwise Prior Authorization/Preservice is required.**

**Bleeding Esophageal Varices**

**Covered Doses:**

Sandostatin IV: Up to 1200 mcg/day

Sandostatin LAR IM: Up to 40 mg every 4 weeks

**ICD-10:**

I85.01-I85.11

**Chemotherapy-Induced Diarrhea, treatment**

**Covered Doses:**

octreotide (Sandostatin IV or LAR Depot)

Sandostatin IV: up to 6000 mcg/ day  
Sandostatin LAR: up to 40 mg every 4 weeks

**ICD-10:**

Encounter Code for Chemotherapy Z51.11 + Diarrhea K52.2, K52.89, or R19.7; OR J9XXX + K52.2, K52.89, or R19.7

**Cryptosporidiosis**

**Covered Doses:**

Sandostatin IV: Up to 2400 mcg/day

**ICD-10:**

A07.2

**Dumping Syndrome**

**Covered Doses:**

Sandostatin IV: Up to 600 mcg/day

Sandostatin LAR IM: Up to 20 mg every 4 weeks

**ICD-10:**

K91.1

**Lymphorrhagia**

**Covered Doses:**

Sandostatin IV: up to 300 mcg/ day

**ICD-10:**

R59

**Malignant Intestinal Obstruction**

**Covered Doses:**

Sandostatin IV: up to 900 mcg per day

Sandostatin LAR: up to 40 mg every 4 weeks

**ICD-10:**

K50.012-K56.69

**Pancreatitis, necrotizing**

**Covered Doses:**

Sandostatin IV: up to 1200 mcg/ day

**ICD-10:**

K85.91, K85.92

**Pituitary Adenomas (TSH-Secreting)**

octreotide (Sandostatin IV or LAR Depot)

**Covered Doses:**

Sandostatin IV: up to 1500 mcg/ day

Sandostatin LAR IM: up to 40 mg every 4 weeks

**ICD-10:**

E23.6

**Polycystic Ovary Syndrome (PCOS)****Covered Doses:**

Sandostatin IV: up to 300 mcg/ day

**ICD-10:**

E28.2

**Radiation-Induced Diarrhea****Covered Doses:**

Sandostatin IV: up to 600 mcg/ day

Sandostatin LAR IM: up to 40 mg intramuscularly every 4 weeks

**ICD-10:**

Encounter for radiotherapy Z51.0, with Diarrhea: K52.2, K52.89, R19.7 OR Effects of radiation, unspecified: T66XXA

**Zollinger-Ellison Syndrome (Gastrinoma)****Covered Doses:**

Sandostatin IV: up to 2000 mcg/day

Sandostatin LAR IM: Up to 60 mg every 4 weeks

**ICD-10:**

D3A.092, E16.4

**References**

1. AHFS. Available by subscription at <http://www.lexi.com>
2. DrugDex. Available by subscription at <http://www.micromedexsolutions.com/home/dispatch>
3. National Comprehensive Cancer Network. Neuroendocrine and Adrenal Tumors (Version 2.2025). Available at <http://www.nccn.org>.
4. National Comprehensive Cancer Network. Palliative Care (Version 2.2025). Available at <http://www.nccn.org>.
5. Scarpellini E, Arts J, Karamanolis G, et al. International consensus on the diagnosis and management of dumping syndrome. Nat Rev Endocrinol. 2020 Aug;16(8):448-466. doi: 10.1038/s41574-020-0357-5. Epub 2020 May 26.
6. National Institute of Diabetes and Digestive and Kidney Diseases. Treatment of Dumping Syndrome. <https://www.niddk.nih.gov/health-information/digestive-diseases/dumping-syndrome/treatment>. Accessed August 15, 2025.

7. Sandostatin LAR Depot (octreotide injection suspension) [prescribing information]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; July 2024.
8. Sandostatin (octreotide injection solution) [prescribing information]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; July 2024.
9. Yavuz MN, Yavuz AA, Aydin F, et al: The efficacy of octreotide in the therapy of acute radiation-induced diarrhea: a randomized controlled study. Int J Radiat Oncol Biol Phys 2002; 54(1):195-202.

#### Review History

Date of Last Annual Review: 4Q2025

Changes from previous policy version:

- No clinical changes following annual review
- For oncology-related indications, medical necessity criteria can be found here: [Blue Shield Oncology-Related Medication Policies](#).
- For PPO, Direct Contract HMO, and when applicable, ASO, and Shared Advantage: Please access Evolent's [CarePro Provider Portal](#) to submit your request.

*Blue Shield of California Medication Policy to Determine Medical Necessity  
Reviewed by P&T Committee*