



Fax completed form to: (855) 840-1678

If this is an URGENT request, please call (800) 882-4462 (800.88.CIGNA)

Sandostatin, Sandostatin LAR Depot

(octreotide LAR Depot, octreotide immediate release)

PHYSICIAN INFORMATION			PATIENT INFORMATION		
* Physician Name:			*Due to privacy regulations we will not be able to respond via fax with the outcome of our review unless all asterisked (*) items on this form are completed.*		
Specialty:	* DEA, NPI or TIN:				
Office Contact Person:			* Patient Name:		
Office Phone:			* Cigna ID:	* Date of Birth:	
Office Fax:			* Patient Street Address:		
Office Street Address:			City:	State:	Zip:
City:	State:	Zip:	Patient Phone:		

Urgency: Standard Urgent (In checking this box, I attest to the fact that applying the standard review time frame may seriously jeopardize the customer's life, health, or ability to regain maximum function)**Medication requested: (please specify name, strength, and dosing schedule)**

ICD10:

- | | | |
|--|---|---|
| <input type="checkbox"/> Octreotide 1000mcg/5mL vial | <input type="checkbox"/> Octreotide 500mcg/mL syringe | <input type="checkbox"/> Octreotide 5000mcg/5mL vial |
| <input type="checkbox"/> Octreotide 500mcg/mL vial | <input type="checkbox"/> Octreotide 0.05mg/mL vial | <input type="checkbox"/> Octreotide 100mcg/mL syringe |
| <input type="checkbox"/> Octreotide 100mcg/mL vial | <input type="checkbox"/> Octreotide 200mcg/mL vial | <input type="checkbox"/> Octreotide 50mcg/mL syringe |
| <input type="checkbox"/> Octreotide 50mcg/mL vial | <input type="checkbox"/> Sandostatin 0.05mg/mL ampule | <input type="checkbox"/> Sandostatin 0.1mg/mL ampule |
| <input type="checkbox"/> Sandostatin 0.5mg/mL ampule | <input type="checkbox"/> Sandostatin LAR Depot 10mg | <input type="checkbox"/> Sandostatin LAR Depot 20mg |
| <input type="checkbox"/> Sandostatin LAR Depot 30mg | | |

Strength and Dosing:

Is this a new start or continuation of therapy**? new start of therapy Continuation of therapy- start date:

If your patient has already begun treatment with drug samples, please choose "new start of therapy". OR if patient has had a break in therapy and is restarting, please choose "new start of therapy".

Where will this medication be obtained?

- | | |
|---|---|
| <input type="checkbox"/> Accredo Specialty Pharmacy** | <input type="checkbox"/> Ambulatory Infusion Center |
| <input type="checkbox"/> Physician's office stock | <input type="checkbox"/> Hospital - In patient |
| <input type="checkbox"/> Home Health / Home Infusion vendor (name): | <input type="checkbox"/> Hospital - Out patient |
| CPT Code(s): _____ | <input type="checkbox"/> Other (please specify): |

**Cigna's nationally preferred specialty pharmacy

Facility and/or doctor dispensing and administering medication:

Facility Name:

State:

Tax ID#:

Address (City, State, Zip Code):

Is this infusion occurring in a facility affiliated with hospital outpatient setting?

Yes No

If yes- Is this patient a candidate for re-direction to an alternate setting after 1-2 infusions (such as AIS, MDO, home) with assistance of a Specialty Care Option Case Manager?

Yes No

NOTE: Per some Cigna plans, infusion of medication MUST occur in the lowest cost, medically appropriate setting.

Is the requested medication for a chronic or long-term condition for which the prescription medication may be necessary for the life of the patient?

 Yes No

Please indicate the condition octreotide, Sandostatin, or Sandostatin LAR is being used to treat and answer additional questions as necessary.

Diagnosis

- Acromegaly
 Enterocutaneous Fistulas
 Meningioma
 Neuroendocrine Tumor(s) [NETs] of the Gastrointestinal Tract, Lung, Thymus (Carcinoid Tumors), and Pancreas (including glucagonomas, gastrinomas, vasoactive intestinal peptides-secreting tumors [VIPomas], insulinomas)
 Pancreatic Fistulas
 Pheochromocytoma and Paraganglioma
 Thymoma and Thymic Carcinoma
 None of the above

(if none of the above) Please provide the patient's diagnosis or reason for treatment.

Clinical Information

(if acromegaly) Did the patient have an inadequate response to surgery and/or radiotherapy? Yes No

(if no) Are surgery and/or radiotherapy NOT an option for this patient? Yes No

(if no) Is the patient experiencing negative effects due to tumor size (for example, optic nerve compression)?
 Yes No

(if acromegaly) Prior to starting any somatostatin analog (for example, Mycapssa [octreotide delayed-release capsules], an octreotide acetate injection product [for example, Bynfezia Pen, Sandostatin {generic}, Sandostatin LAR Depot], Signifor LAR [pasireotide injection], Somatuline Depot [lanreotide injection], dopamine agonist [for example, cabergoline, bromocriptine], or Somavert [pegvisomant injection]), does/did your patient have an insulin-like growth factor-1 (IGF-1) level above the upper limit of normal based on age and gender for the reporting laboratory (note that references ranges for IGF-1 vary among laboratories)? Yes No

(if acromegaly) Is the requested medication being prescribed by (or in consultation with) an endocrinologist? Yes No

(if Meningioma) Is the requested medication being prescribed by (or in consultation with) an oncologist, radiologist, or neurosurgeon?
 Yes No

(if NETs) Is the requested medication being prescribed by (or in consultation with) an oncologist, endocrinologist, or gastroenterologist?
 Yes No

(if Pancreatic Fistulas) Is the patient being treated for operative trauma, pancreatic resection, acute or chronic pancreatitis, or pancreatic infection?
 Yes No

(if Pheochromocytoma and Paraganglioma) Is the requested medication being prescribed by (or in consultation with) an endocrinologist, oncologist, or neurologist?
 Yes No

(if Thymoma and Thymic Carcinoma) Is the requested medication being prescribed by (or in consultation with) an oncologist?
 Yes No

If requesting Sandostatin LAR Depot, preferred product criteria must be met:

(if acromegaly) Has it been documented that the patient has tried Somatuline Depot? Yes No

(if NETs) Has it been documented that the patient has tried Somatuline Depot subcutaneous injection? Yes No

(if no) Has the patient already been started on therapy with Sandostatin LAR? Yes No

(if Pheochromocytoma and Paraganglioma) Has it been documented that the patient has tried Somatuline Depot? Yes No

(if no) Has the patient already been started on therapy with Sandostatin LAR? Yes No

Additional pertinent information: *(please include clinical reasons for drug, relevant lab values, etc.)*

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature: _____ **Date:** _____

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Our standard response time for prescription drug coverage requests is 5 business days. If your request is urgent, it is important that you call us to expedite the request. View our Prescription Drug List and Coverage Policies online at cigna.com.

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