

Clinical Policy: Lutetium Lu 177 Dotatate (Lutathera)

Reference Number: LA.PHAR.384

Effective Date: 05.22.18

Last Review Date: 02.23

Line of Business: Medicaid

[Coding Implications](#)

[Revision Log](#)

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

*****Please note: This policy is for medical benefit*****

Description

Lutetium Lu 177 dotatate (Lutathera®) is a radiolabeled somatostatin analog.

FDA Approved Indication(s)

Lutathera is indicated for the treatment of somatostatin receptor-positive gastroenteropancreatic neuroendocrine tumors (GEP-NETs), including foregut, midgut, and hindgut NETs in adults.

Policy/Criteria

Prior authorization is required. Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

It is the policy of Louisiana Healthcare Connections that Lutathera is **medically necessary** when the following criteria are met:

I. Initial Approval Criteria

A. Neuroendocrine Tumors (must meet all):

1. Diagnosis of a somatostatin receptor-positive NET of one of the following origins (a or b):
 - a. Gastrointestinal tract or pancreas;
 - b. Lung or thymus (off-label);
2. Prescribed by or in consultation with an oncologist;
3. Age \geq 18 years;
4. Disease is metastatic or locally advanced, and unresectable;
5. Member experienced disease progression while on a somatostatin analog (e.g., octreotide, lanreotide);
6. Dose does not exceed 7.4 GBq (200 mCi) every 8 weeks, up to a total of 4 doses.

Approval duration: 32 weeks (no more than 4 total doses)

B. Pheochromocytoma/Paraganglioma (off-label) (must meet all):

1. Diagnosis of a somatostatin receptor-positive pheochromocytoma/paraganglioma;
2. Prescribed by or in consultation with an oncologist;
3. Disease is metastatic or locally advanced, and unresectable;
4. Dose does not exceed 7.4 GBq (200 mCi) every 8 weeks, up to a total of 4 doses.

Approval duration: 32 weeks (no more than 4 total doses)

C. Other diagnoses/indications

1. Refer to the off-label use policy if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): LA.PMN.53 for Medicaid.

II. Continued Therapy

A. All Indications in Section I (must meet all):

1. Currently receiving medication via Louisiana Healthcare Connections benefit, or documentation supports that member is currently receiving Lutathera for a covered indication;
2. Member is responding positively to therapy;
3. Member has not received ≥ 4 doses of Lutathera;
4. If request is for a dose increase, new dose does not exceed 7.4 GBq (200 mCi) every 8 weeks, up to a total of 4 doses.

Approval duration: 32 weeks (no more than 4 total doses)

B. Other diagnoses/indications (must meet 1 or 2):

1. Currently receiving medication via Louisiana Healthcare Connections benefit and documentation supports positive response to therapy.
Approval duration: Duration of request or 6 months (whichever is less); or
2. Refer to the off-label use policy if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): LA.PMN.53 for Medicaid.

III. Diagnoses/Indications for which coverage is NOT authorized:

- A. Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policy – LA.PMN.53 for Medicaid.

IV. Appendices/General Information

Appendix A: Abbreviation/Acronym Key

CT: computed tomography	mCi: millicurie
FDA: Food and Drug Administration	NCCN: National Comprehensive Cancer Network
GEP-NET: gastroenteropancreatic neuroendocrine tumor	

Appendix B: Therapeutic Alternatives

This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may require prior authorization.

Drug Name	Dosing Regimen	Dose Limit/ Maximum Dose
Somatuline [®] Depot (lanreotide)	90 – 120 mg SC every 4 weeks	120 mg/month
Sandostatin [®] LAR Depot (octreotide LAR)*	20 – 30 mg IM once monthly (20 mg may be used for pancreatic NETs)	30 mg/month

Drug Name	Dosing Regimen	Dose Limit/ Maximum Dose
Sandostatin® (octreotide)	150 – 250 mcg SC TID	450 mcg/day

Therapeutic alternatives are listed as Brand name® (generic) when the drug is available by brand name only and generic (Brand name®) when the drug is available by both brand and generic.

**Off-label for the treatment of NETs (octreotide is only FDA-approved for the treatment of symptoms associated with carcinoid tumors) – NET dosing recommendations are per the NCCN guidelines*

Appendix C: Contraindications/Boxed Warnings

None reported

Appendix D: General Information

- Somatostatin receptor expression can be detected by somatostatin receptor-based imaging, which includes ⁶⁸Ga-dotatate PET/CT (preferred per the NCCN) and somatostatin receptor scintigraphy.
- Use of Lutathera with somatostatin analogs:
 - Before initiating Lutathera: Long-acting somatostatin analogs (e.g., long-acting octreotide) should be discontinued for at least 4-6 weeks prior to initiation of Lutathera. Short-acting octreotide can be administered as needed up to 24 hours prior to initiating Lutathera.
 - After Lutathera: Administer long-acting octreotide 30 mg intramuscularly 4 to 24 hours after each Lutathera dose and short-acting octreotide for symptomatic management.
 - Continue long-acting octreotide 30 mg intramuscularly every 4 weeks after completing Lutathera until disease progression or for up to 18 months following treatment initiation.
 - During Lutathera treatment: IV infusion of amino acids is critical for nephron protection and should be infused 30 minutes before and 3 hours after Lutathera treatment
 - Following Lutathera treatment: Octreotide or lanreotide (short and long acting) can be administered 4 to 24 hours after completing Lutathera.

V. Dosage and Administration

Indication	Dosing Regimen	Maximum Dose
GEP-NET	7.4 GBq (200 mCi) IV every 8 weeks for a total of 4 doses	7.4 BGq (200 mCi) IV (4 doses)
NET of lung or thymus origin, pheochromocytoma, paraganglioma*		

**Off-label – dosing recommendations are per the NCCN guidelines*

VI. Product Availability

Single-dose vial for injection: 370 MBq/mL (10 mCi/mL)

VII. References

1. Lutathera Prescribing Information. Millburn, NJ: Advanced Accelerator Applications USA, Inc.; June 2021. Available at: <https://www.lutathera.com>. Accessed May 3, 2022.
2. National Comprehensive Cancer Network. Neuroendocrine and Adrenal Tumors. Version 4.2021. Available at:

- https://www.nccn.org/professionals/physician_gls/pdf/neuroendocrine.pdf. Accessed May 3, 2022.
3. National Comprehensive Cancer Network Drugs and Biologics Compendium. Available at: http://www.nccn.org/professionals/drug_compendium. Accessed May 3, 2022.
 4. Strosberg J, El-Haddad G, Wolin E, et al. Phase 3 trial of ¹⁷⁷Lu-dotatate for midgut neuroendocrine tumors. *N Engl J Med*. 2017; 376(2): 125-135.
 5. Brabander T, van der Zwan WA, Teunissen JJM, et al. Long-term efficacy, survival, and safety of [¹⁷⁷Lu-DOTA⁰,Tyr³]octreotate in patients with gastroenteropancreatic and bronchial neuroendocrine tumors. *Clin Cancer Res*. 2017; 1-8.
 6. Clinical Pharmacology [database online]. Elsevier, Inc.; 2022. Available at: <https://www.clinicalkey.com/pharmacology/>.

Coding Implications

Codes referenced in this clinical policy are for informational purposes only. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

HCP Codes	Description
A9513	Lutetium Lu 177, dotatate, therapeutic, 1 millicurie (mCi)

Reviews, Revisions, and Approvals	Date	LDH Approval Date
Converted corporate to local policy	02.23	

Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. LHCC makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved.

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