SENTARA COMMUNITY PLAN (MEDICAID)

MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions:</u> The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; <u>fax to 1-844-305-2331</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If information provided is not complete, correct, or legible, authorization can be delayed.</u>

Drug Requested: Lutathera® (lutetium Lu 177 dotatate) IV (A9513)

MEMBER & PRESCRIBER INFO	ORMATION: Authorization may be delayed if incomplete.
Member Name:	
Member Sentara #:	Date of Birth:
Prescriber Name:	
Prescriber Signature:	Date:
Office Contact Name:	
Phone Number:	Fax Number:
DEA OR NPI #:	
DRUG INFORMATION: Authorize	ation may be delayed if incomplete.
Drug Form/Strength:	
Dosing Schedule:	Length of Therapy:
Diagnosis:	ICD Code, if applicable:
Weight:	Date:
☐ Standard Review. In checking this box.	, the timeframe does not jeopardize the life or health of the member num function and would not subject the member to severe pain.

Quantity Limits:

- A. Length of Authorization
 - Coverage will be provided for 1 year (4 doses only) and may **NOT** be renewed.
- B. Max Units (per dose and over time) [HCPCS Unit]:
 - 200 billable units (7.4 GBq = 200 mCi) every 8 weeks for a total of 4 doses

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

Approval Criteria – Coverage cannot be renewed

☐ Member is at least 12 years of age or older

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	Requesting provider is an oncologist		
	For female members of reproductive potential, a negative pregnancy test has been confirmed		
		nber has progressive locally advanced or metastatic, somatostatin receptor-positive roenteropancreatic neuroendocrine tumors (GEP-NETs)	
	Member's disease is somatostatin receptor-positive in all tumor lesions (OctreoScan uptake \geq normal liver)		
	Member has well-differentiated disease with a Ki67 labeling index score of $\leq 20\%$		
	Member's condition has progressed on long-acting octreotide or lanreotide		
	Member will discontinue any long-acting somatostatin analogues (e.g., octreotide LAR, pasireotide LAR, lanreotide depot) within the previous 4 weeks OR short-acting somatostatin analogues (e.g., octreotide, pasireotide) within 24 hours prior to therapy		
	Requested medication will be used in combination with a long-acting somatostatin analog (e.g., octreotide LAR, lanreotide depot) given as a single-injection (between 4-24 hours) following each Lutathera infusion		
	NOTE: Somatostatin analogs require separate prior authorization		
	Provider will follow the recommended dosage per weight and timeline indication detailed in the table below:		
ndi	cation	Dose	
euroendocrine tumors GEP-NETs)		• 7.4 GBq (200 mCi) every 8 weeks for a total of 4 doses. Administer a single dose of long-acting somatostatin analog between 4 to 24 hours after each Lutathera dose. (Long-acting somatostatin analog may not be repeated until	

Indication	Dose
Gastroenteropancreatic neuroendocrine tumors (GEP-NETs)	 7.4 GBq (200 mCi) every 8 weeks for a total of 4 doses. Administer a single dose of long-acting somatostatin analog between 4 to 24 hours after each Lutathera dose. (Long-acting somatostatin analog may not be repeated until after the next scheduled dose of Lutathera to provide the 4-week drug-free interval. Short-acting octreotide may be administered up to 24 hours prior to each Lutathera dose) Initiate recommended intravenous amino acid solution 30 minutes before
	each Lutathera dose)

Reauthorization Criteria – Coverage cannot be renewed

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Medication being provided by (check box below that applies):
□ Location/site of drug administration:
NPI or DEA # of administering location:
<u>OR</u>
□ Specialty Pharmacy – Proprium Rx
For urgent reviews: Practitioner should call Sentara Health Pre-Authorization Department if they believe a standard review would subject the member to adverse health consequences. Sentara Health's definition of urgent is a lack of treatment that could seriously jeopardize the life or health of the member or the member's ability to regain maximum function.
Use of samples to initiate therapy does not meet step edit/ preauthorization criteria. *Previous therapies will be verified through pharmacy paid claims or submitted chart notes.*