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 (<u>including phone and fax #s</u>) on this form is correct. <u>If information provided is not complete</u>, correct, or legible, authorization can be delayed.

Drug Requested: Pluvicto® (lutetium Lu 177 vipivotide tetraxetan) IV (A9699) (Medical)

Member Name:	
Member Sentara #:	Date of Birth:
Prescriber Name:	
Prescriber Signature:	
Office Contact Name:	
Phone Number:	
DEA OR NPI #:	
DRUG INFORMATION: Authoriz	
Drug Form/Strength:	
Dosing Schedule:	Length of Therapy:
Diagnosis:	ICD Code:
Weight:	Date:

Quantity Limits:

A. Length of Authorization:

- Coverage will be provided for six months (4 doses) and may be renewed to provide for 2 additional doses (3-months)
- The total number of doses authorized cannot exceed 6 doses

B. Max Units (per dose and over time) [HCPCS Unit]:

- 200 mCi (7.4 GBq = 200 mCi) every 6 weeks for a total of 6 doses
- Pluvicto 1,000 MBq/mL (27 mCi/mL) of lutetium Lu 177 vipivotide tetraxetan: 30 mL single- dose vial containing 7.4 GBq (200 mCi)

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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.

Initial Authorization: 4 doses

- ☐ Member is at least 18 years of age
- ☐ Requesting provider is an oncologist
- ☐ Member has a diagnosis of metastatic castration-resistant prostate cancer (mCRPC)
- ☐ Member's disease is confirmed to be prostate-specific membrane antigen (PSMA)-positive [defined as Ga-68 gozetotide uptake greater than that of liver parenchyma in one or more metastatic lesions of any size in any organ system]
- ☐ Member will receive concurrent treatment with a gonadotropin releasing hormone (GnRH)-analog, **OR** has had a bilateral orchiectomy
- ☐ Member has been previously treated with an androgen receptor pathway inhibitor (e.g., enzalutamide, abiraterone) **AND** taxane-based chemotherapy (e.g., docetaxel)
- ☐ Provider will follow the recommended dosage per weight and timeline indication detailed in the table below:

Indication	Dose
mCRPC	• The recommended Pluvicto dosage is 7.4 GBq (200 mCi) intravenously every 6 weeks (up to 10 weeks for toxicities) for up to 6 doses, or until disease progression, or unacceptable toxicity

Reauthorization: Additional 2 doses. 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.

- ☐ Confirmation of disease response with initial treatment as defined by stabilization of disease or at least a partial response has been documented and submitted by provider
- ☐ Member has experienced an absence of unacceptable toxicity from the drug (e.g., bone marrow suppression, nephrotoxicity)
- ☐ Member has <u>NOT</u> received more than 6 total doses

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Medication being provided by (check box below that applies):	
□ Location/site of drug administration:	
NPI or DEA # of administering location:	
OR	
□ Specialty Pharmacy - PropriumRx	

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.