

Member Name: _____ Date of Birth: _____ Member ID#: _____

Drug Information

Prescriber billing (HCPCS code: _____) Start Date (or date of next dose): _____

Dose: _____ Regimen: _____

Billing Provider Information

Provider NPI: _____ Provider Name: _____

Provider Phone: _____ Provider Fax: _____

Prescriber Information

Prescriber NPI: _____ Prescriber Name: _____

Prescriber Phone: _____ Prescriber Fax: _____ Specialty: _____

Criteria

For Initial Authorization:

1. Please indicate the diagnosis and information:

Gastroenteropancreatic Neuroendocrine (GEP-NET)

A. Is diagnosis progressive locoregional advanced disease or metastatic disease? Yes ___ No ___

B. Is there positive imaging of somatostatin receptors? Yes ___ No ___

C. Will Lutathera® be used as second-line or subsequent therapy following progression on octreotide or lanreotide? Yes ___ No ___

D. Will Lutathera® be used as first-line for treatment of pheochromocytoma/paraganglioma? Yes ___ No ___

If diagnosis is not listed above, please indicate diagnosis: _____

Additional Information: _____

For Continued Authorization:

1. Date of last dose: _____

2. Does member have any evidence of progressive disease while on Lutathera®? Yes ___ No ___

3. Has the member experienced any adverse drug reactions related to Lutathera® therapy? Yes ___ No ___

If yes, please specify adverse reactions: _____

Additional Information: _____

Prescriber Signature: _____ Date: _____

I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge. Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy
Pharmacy Management Consultants
Product Based Prior Authorization Unit
Fax: 1-800-224-4014
Phone: 1-800-522-0114 Option 4

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