

UNIVERSITY OF MISSOURI  
RADIOPHARMACEUTICAL ADMINISTRATION FORM  
Lu-177 Pluvicto ONLY

**This form shall be used for the administration of Therapeutic dosages of *Lu-177 Pluvicto* only**

**RADIOPHARMACEUTICAL PRESCRIPTION:  
Lutetium-177 PSMA (Pluvicto)**

(To be completed by authorized physician)

**Patient sticker here**

To include:

- Patient Name
- DOB
- Hospital ID number

Procedure: *Lutetium-177 PSMA for Prostate Cancer*

Requesting Physician: \_\_\_\_\_

Authorizing Physician: \_\_\_\_\_

Radiopharmaceutical Prescribed: \_\_\_\_\_

Route of administration: *IV Administration*

Dosage Ordered: \_\_\_\_\_ mCi

Infusion Cycle Number: \_\_\_\_\_

Signature of Authorized Physician: \_\_\_\_\_

Date \_\_\_\_\_

Written instructions provided.

**PRE-ADMINISTRATION**

Prior to administering the radiopharmaceutical, the person performing the administration must verify the identity of the patient in the written directive, and that the details of the administration are in accordance with the written directive and approved by Nuclear Medicine/Radiation Oncology consult. The person responsible for the administration of the radiopharmaceutical will review relevant labs to determine that they are satisfactory and complete the form.

**PATIENT IDENTIFICATION VERIFIED** by (2 required):

- Name                                       Hospital I.D.  
 Date of Birth    Personal Recognition    Photo I.D.

Patient is:

- Male

***VERIFY PATIENT:***

- Patient has consented  
**AND**  
 Patient had PSMA Scan for Prostate Ca.  
 Metastatic disease by PSMA Scan  
 Satisfactory blood counts.

**RADIOPHARMACEUTICAL DOSAGE VERIFICATION:** (COMPLETED BY PERSON ADMINISTERING RADIOPHARMACEUTICAL)

Radiopharmaceutical drug being administered: Lutetium-177 Pluvicto

Radiopharmaceutical Lot Number: \_\_\_\_\_

Pharmaceutical Company measurement (Dose Received): \_\_\_\_\_ mCi Date: \_\_\_\_/\_\_\_\_/\_\_\_\_ Time: \_\_\_\_:\_\_\_\_

MU Dose Calibrator measurement (Dose Prepared): [KK] \_\_\_\_\_ mCi Date: \_\_\_\_/\_\_\_\_/\_\_\_\_ Time: \_\_\_\_:\_\_\_\_

Route of Administration: IV Administration

Pre-/Post-Administration checks performed by signature: \_\_\_\_\_ Date: \_\_\_\_/\_\_\_\_/\_\_\_\_ Time: \_\_\_\_:\_\_\_\_

**RADIOPHARMACEUTICAL INFUSION:**

Signature of Individual Administering Dose: \_\_\_\_\_ Date: \_\_\_\_/\_\_\_\_/\_\_\_\_ Time: \_\_\_\_:\_\_\_\_

IV SITE: \_\_\_\_\_  Patency Assessment performed

Pluvicto Infusion Start Time: \_\_\_\_:\_\_\_\_ Pluvicto Infusion Completion time: \_\_\_\_:\_\_\_\_

**POST-INFUSION:**

Note: Revisions must be made, signed, and dated by an AU within 24 hours

Calculated Activity Delivered [DDD]: \_\_\_\_\_ mCi Percent Prescribed Activity Delivered [EEE] \_\_\_\_\_ %

Emerging patient conditions?  Yes  No

Provide explanation:

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

Authorized User Signature: \_\_\_\_\_ Date: \_\_\_\_/\_\_\_\_/\_\_\_\_

Comments/Notes: \_\_\_\_\_

**Patient Released by the Following Criteria**

Patient reading at 1 m is <8.6mR/hr.

Patient release instructions given

Released by NM (Signature): \_\_\_\_\_ Date: \_\_\_\_/\_\_\_\_/\_\_\_\_

Final review by Nuclear Medical Staff (Signature): \_\_\_\_\_ Date: \_\_\_\_/\_\_\_\_/\_\_\_\_

RETAIN THIS RECORD FOR 3 YEARS: REQUIRED BY 10 CFR 35.2040