

Quadramet® (Samarium Sm 153 Lexidronam) (Intravenous)

Document Number: OHSU HEALTHSERVICES-0435

Last Review Date: 07/05/2022

Date of Origin: 03/04/2019

Dates Reviewed: 03/2019, 07/2020, 07/2021, 07/2022

I. Length of Authorization

Coverage will be provided for 1 treatment course and may be renewed, one-time only, after 60 days.

II. Dosing Limits

A. Quantity Limit (max daily dose) [NDC Unit]:

- N/A

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

- 1 billable unit (up to 150 mCi) per treatment course

III. Initial Approval Criteria ¹

Coverage is provided in the following conditions:

- Patient is at least 16 years of age; **AND**
- Women of child-bearing age must have a negative pregnancy test prior to treatment; **AND**
- Lactating women should discontinue breast feeding at least 6 weeks prior to administration; **AND**

Universal Criteria ¹

- Patient will not use in combination with or has not had a treatment course of strontium-89 chloride within the previous 90 days; **AND**
- Patient has not had a treatment course of samarium-sm-153 lexidronam within the previous 60 days; **AND**
- Patients of reproductive potential will use effective contraception during treatment with therapy and for at least 6 months after the last dose; **AND**
- Patient does not have significant bone marrow suppression (i.e., neutropenia, leukopenia, thrombocytopenia, etc.); **AND**

- Patient does not have disseminated intravascular coagulation; **AND**

Pain related to metastatic bone lesions † 1,2

- Used for palliative treatment of metastatic skeletal bone pain; **AND**
- Patient has had a positive (enhancement) radionuclide bone scan confirming osteoblastic metastatic bone lesions; **AND**
- Therapy will not be used for spinal cord compression pain; **AND**
- Patient has failed other conventional treatments for bone pain due to skeletal metastases (e.g., chemotherapy, hormonal therapy, external beam radiation, opioid analgesics, etc.); **AND**
- Patient has a life-expectancy of at least 6 months

Osteosarcoma ‡ 5,7

- Used for relapsed or refractory disease beyond second-line therapy

† FDA Approved Indication(s); ‡ Compendia recommended indication(s); Φ Orphan Drug

IV. Renewal Criteria 1,4,5

Coverage can be renewed based upon the following criteria:

- Patient continues to meet universal and other indication specific relevant criteria such as concomitant therapy requirements (not including prerequisite therapy), performance status, etc. identified in section III; **AND**
- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include: severe leukopenia, severe thrombocytopenia, severe neutropenia, etc.; **AND**
- Patient has experienced hematological recovery since administration of the initial dose; **AND**

Pain related to metastatic bone lesions

- Patient had an inadequate response or recurrence of bone pain after the initial dose

Osteosarcoma

- Disease response with treatment as defined by stabilization of disease or decrease in size of tumor or tumor spread

V. Dosage/Administration 1,7

Indication	Dose
Metastatic Bone Pain	The recommended dose is 1.0 mCi/kg, administered intravenously over a period of one minute through a secure in-dwelling catheter and followed with a saline flush.

	<ul style="list-style-type: none"> – The patient should ingest (or receive by IV administration) a minimum of 500 mL (2 cups) of fluids prior to injection and should void as often as possible after injection to minimize radiation exposure to the bladder.
Osteosarcoma	1.21 mCi/kg administered intravenously
<p>– <i>Quadramet is a radiopharmaceutical; handle with appropriate safety measures to minimize radiation exposure. Use waterproof gloves and effective radiation shielding when handling. Quadramet should be used by or under the control of physicians who are qualified by specific training and experience in the safe use and handling of radiopharmaceuticals, and whose experience and training have been approved by the appropriate governmental agency authorized to license the use of radiopharmaceuticals.</i></p>	
<p>– <i>Thaw at room temperature before administration and use within 8 hours of thawing.</i></p>	

VI. Billing Code/Availability Information

HCPCS Code:

- A9604 – Samarium sm-153 lexidronam, therapeutic, per treatment dose, up to 150 millicuries

NDC:

- Quadramet 5550 MBq (150 mCi) 3 mL frozen single-dose vial: 11994-0016-XX

VII. References

1. Quadramet [package insert]. N. Billerica, MA; Lantheus Medical Imaging, Inc.; September 2017. Accessed June 2022.
2. Anderson PM, Wiseman GA, Dispenzieri A, et al. High-dose samarium-153 ethylene diamine tetramethylene phosphonate: low toxicity of skeletal irradiation in patients with osteosarcoma and bone metastases. *J Clin Oncol.* 2002 Jan 1;20(1):189-96.
3. American College of Radiology (ACR), American Society for Radiation Oncology (ASTRO). ACR-ASTRO practice guideline for the performance of therapy with unsealed radiopharmaceutical sources. [online publication]. Reston, VA: American College of Radiology (ACR); 2011.
4. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) Bone Cancer. Version 2.2022. National Comprehensive Cancer Network, 2022. The NCCN Compendium® is a derivative work of the NCCN Guidelines®. NATIONAL COMPREHENSIVE CANCER NETWORK®, NCCN®, and NCCN GUIDELINES® are trademarks owned by the National Comprehensive Cancer Network, Inc. To view the most recent and complete version of the Compendium, go online to NCCN.org. Accessed June 2022.
5. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) Samarium Sm-153 EDTMP. National Comprehensive Cancer Network, 2022. The NCCN Compendium® is a derivative work of the NCCN Guidelines®. NATIONAL COMPREHENSIVE CANCER NETWORK®, NCCN®, and NCCN GUIDELINES® are trademarks owned by the National Comprehensive Cancer Network, Inc. To view the most recent and complete version of the Compendium, go online to NCCN.org. Accessed June 2022.

6. American College of Radiology (ACR), American College of Nuclear Medicine (ACNM), American Society for Radiation Oncology (ASTRO), and Society of Nuclear Medicine and Molecular Imaging (SNMMI). ACR–ACNM–ASTRO–SNMMI Practice Parameter for the Performance of Therapy with Unsealed Radiopharmaceutical Sources. [online publication]. Reston, VA: American College of Radiology (ACR); Revised 2019 (Resolution 41). <https://www.acr.org/-/media/ACR/Files/Practice-Parameters/unsealedsources.pdf>
7. Loeb DM, Garrett-Mayer E, Hobbs RF, et al. Dose-finding study of ¹⁵³Sm-EDTMP in patients with poor-prognosis osteosarcoma. *Cancer*. 2009;115(11):2514-2522. doi:10.1002/cncr.24286

Appendix 1 – Covered Diagnosis Codes

ICD10	ICD-10 Description
C79.51	Secondary malignant neoplasm of bone
C79.52	Secondary malignant neoplasm of bone marrow
C40.00	Malignant neoplasm of scapula and long bones of unspecified upper limb
C40.01	Malignant neoplasm of scapula and long bones of right upper limb
C40.02	Malignant neoplasm of scapula and long bones of left upper limb
C40.10	Malignant neoplasm of short bones of unspecified upper limb
C40.11	Malignant neoplasm of short bones of right upper limb
C40.12	Malignant neoplasm of short bones of left upper limb
C40.20	Malignant neoplasm of long bones of unspecified lower limb
C40.21	Malignant neoplasm of long bones of right lower limb
C40.22	Malignant neoplasm of long bones of left lower limb
C40.30	Malignant neoplasm of short bones of unspecified lower limb
C40.31	Malignant neoplasm of short bones of right lower limb
C40.32	Malignant neoplasm of short bones of left lower limb
C40.80	Malignant neoplasm of overlapping sites of bone and articular cartilage of unspecified limb
C40.81	Malignant neoplasm of overlapping sites of bone and articular cartilage of right limb
C40.82	Malignant neoplasm of overlapping sites of bone and articular cartilage of left limb
C40.90	Malignant neoplasm of unspecified bones and articular cartilage of unspecified limb
C40.91	Malignant neoplasm of unspecified bones and articular cartilage of right limb
C40.92	Malignant neoplasm of unspecified bones and articular cartilage of left limb
C41.0	Malignant neoplasm of bones of skull and face
C41.1	Malignant neoplasm of mandible
C41.2	Malignant neoplasm of vertebral column
C41.3	Malignant neoplasm of ribs, sternum and clavicle
C41.4	Malignant neoplasm of pelvic bones, sacrum and coccyx
C41.9	Malignant neoplasm of bone and articular cartilage, unspecified
Z85.830	Personal history of malignant neoplasm of bone

Appendix 2 – Centers for Medicare and Medicaid Services (CMS)

Medicare coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determination (NCD), Local Coverage Articles (LCAs) and Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. They can be found at: <https://www.cms.gov/medicare-coverage-database/search.aspx>. Additional indications may be covered at the discretion of the health plan.

Medicare Part B Covered Diagnosis Codes (applicable to existing NCD/LCA/LCD): N/A

Medicare Part B Administrative Contractor (MAC) Jurisdictions		
Jurisdiction	Applicable State/US Territory	Contractor
E (1)	CA, HI, NV, AS, GU, CNMI	Noridian Healthcare Solutions, LLC
F (2 & 3)	AK, WA, OR, ID, ND, SD, MT, WY, UT, AZ	Noridian Healthcare Solutions, LLC
5	KS, NE, IA, MO	Wisconsin Physicians Service Insurance Corp (WPS)
6	MN, WI, IL	National Government Services, Inc. (NGS)
H (4 & 7)	LA, AR, MS, TX, OK, CO, NM	Novitas Solutions, Inc.
8	MI, IN	Wisconsin Physicians Service Insurance Corp (WPS)
N (9)	FL, PR, VI	First Coast Service Options, Inc.
J (10)	TN, GA, AL	Palmetto GBA, LLC
M (11)	NC, SC, WV, VA (excluding below)	Palmetto GBA, LLC
L (12)	DE, MD, PA, NJ, DC (includes Arlington & Fairfax counties and the city of Alexandria in VA)	Novitas Solutions, Inc.
K (13 & 14)	NY, CT, MA, RI, VT, ME, NH	National Government Services, Inc. (NGS)
15	KY, OH	CGS Administrators, LLC