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# Production and Supply of $\alpha$ -Particle–Emitting Radionuclides for Targeted $\alpha$ -Therapy

Valery Radchenko<sup>1,2</sup>, Alfred Morgenstern<sup>3</sup>, Amir R. Jalilian<sup>4</sup>, Caterina F. Ramogida<sup>1,5</sup>, Cathy Cutler<sup>6</sup>, Charlotte Duchemin<sup>7,8</sup>, Cornelia Hoehr<sup>1</sup>, Ferrid Haddad<sup>9</sup>, Frank Bruchertseifer<sup>3</sup>, Haavar Gausemel<sup>10</sup>, Hua Yang<sup>1</sup>, Joao Alberto Osso<sup>4</sup>, Kohshin Washiyama<sup>11</sup>, Kenneth Czerwinski<sup>12</sup>, Kirsten Leufgen<sup>13</sup>, Marek Pruszyński<sup>14,15</sup>, Olga Valzdorf<sup>16</sup>, Patrick Causey<sup>17</sup>, Paul Schaffer<sup>1</sup>, Randy Perron<sup>18</sup>, Samsonov Maxim<sup>19</sup>, D. Scott Wilbur<sup>20</sup>, Thierry Stora<sup>7</sup>, and Yawen Li<sup>20</sup>

<sup>1</sup>Life Sciences Division, TRIUMF, Vancouver, British Columbia, Canada; <sup>2</sup>Department of Chemistry, University of British Columbia, Vancouver, British Columbia, Canada; <sup>3</sup>European Commission, Joint Research Centre, Karlsruhe, Germany; <sup>4</sup>International Atomic Energy Agency, Vienna, Austria; <sup>5</sup>Simon Fraser University, Burnaby, British Columbia, Canada; <sup>6</sup>Collider Accelerator Department, Brookhaven National Laboratory, Upton, New York; <sup>7</sup>CERN, Geneva, Switzerland; <sup>8</sup>Institute for Nuclear and Radiation Physics, KU Leuven, Heverlee, Belgium; <sup>9</sup>Arronax, Nantes, France; <sup>10</sup>Bayer American Samoa, Oslo, Norway; <sup>11</sup>Advanced Clinical Research Center, Fukushima Medical University, Fukushima, Japan; <sup>12</sup>Radiochemistry Program, Department of Chemistry, University of Nevada, Las Vegas, Nevada; <sup>13</sup>SCIPROM Sarl, Saint-Sulpice, Switzerland; <sup>14</sup>Institute of Nuclear Chemistry and Technology, Warsaw, Poland; <sup>15</sup>NOMATEN Centre of Excellence, National Centre for Nuclear Research, Otwock, Poland; <sup>16</sup>Isotope JSC, Rosatom State Corp., Moscow, Russian Federation; <sup>17</sup>BWXT Medical Ltd., Ottawa, Ontario, Canada; <sup>18</sup>Canadian Nuclear Laboratories, Chalk River, Ontario, Canada; <sup>19</sup>State Scientific Centre of the Russian Federation, Lepunsky Institute for Physics and Power Engineering, Rosatom State Corp., Obninsk, Russian Federation; and <sup>20</sup>Department of Radiation Oncology, University of Washington, Seattle, Washington

Encouraging results from targeted  $\alpha$ -therapy have received significant attention from academia and industry. However, the limited availability of suitable radionuclides has hampered widespread translation and application. In the present review, we discuss the most promising candidates for clinical application and the state of the art of their production and supply. In this review, along with 2 forthcoming reviews on chelation and clinical application of  $\alpha$ -emitting radionuclides, *The Journal of Nuclear Medicine* will provide a comprehensive assessment of the field.

**Key Words:** targeted  $\alpha$ -therapy; production and supply of radionuclides; <sup>227</sup>Th/<sup>223</sup>Ra; <sup>225</sup>Ac/<sup>213</sup>Bi; <sup>211</sup>At; <sup>212</sup>Pb/<sup>212</sup>Bi

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**T**argeted radionuclide therapy has seen important clinical breakthroughs, notably originating from the successful clinical translation of prostate-specific membrane antigen–targeted and somatostatin receptor–targeted therapy with  $\beta^-$  emitters (notably <sup>177</sup>Lu) (1,2).  $\alpha$ -emitting radionuclides have also been applied successfully in research and the clinic. Although not a bioconjugate, Xofigo (<sup>223</sup>RaCl<sub>2</sub>; Bayer) received clinical approval, representing an important milestone in the translation and application of  $\alpha$ -emitter–based radiopharmaceuticals (3).

Improved access to a portfolio of selective  $\alpha$ -emitting bioconjugates and radiopharmaceuticals is an important requirement for

preclinical evaluations, clinical trials, and translation (4,5). Targeted  $\alpha$ -therapy (TAT) combines  $\alpha$ -emitting radionuclides with selective delivery systems (e.g., peptides or antibodies). Because of their high linear-energy transfer and high energy (several mega-electronvolts), TAT radiopharmaceuticals deliver therapeutic power within a range of a few cell diameters. This power generates maximal damage to targeted cells while minimizing off-target effects on healthy tissues (6).

Significant efforts are required to optimize the formulation of stable radiopharmaceuticals, determine microdosimetry, and advance clinical studies. However, the major bottleneck for conducting translational research with  $\alpha$ -emitters is their limited availability. The high atomic number (Z) of TAT radionuclides, resulting in complex production and lengthy irradiation using powerful reactors or cyclotrons, creates this problem. Alternatively, irradiation of highly radioactive targets at specialized facilities or generation from uncommon isotopes may be required. Therefore, the demand for  $\alpha$ -emitters often significantly exceeds availability and supply.

Several comprehensive reviews about various aspects of TAT, including radiochemical considerations (7), and preclinical and clinical applications have been published (8,9). In the current review, we asked a group of experts to highlight research challenges and opportunities for the rapidly evolving field of TAT. We describe the state of the art in production and supply of the most potent clinically relevant  $\alpha$ -emitters. We also highlight discrepancies between demand and availability.

Member states have asked the International Atomic Energy Agency to assist with capacity building and technology transfer for the development, production, and quality control of new generations of therapeutic radiopharmaceuticals, including  $\alpha$ -emitters. During technical meetings at the International Atomic Energy Agency in 2013 (10), 2018 (11), and 2019, the demand, production routes, radiopharmaceutical aspects, and supply of <sup>225</sup>Ac were extensively discussed. The International Atomic Energy Agency will provide

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For correspondence or reprints, contact Valery Radchenko (vradchenko@triumf.ca).

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guidelines to the member states for production, quality control, pre-clinical tests, and waste management of  $\alpha$ -radiopharmaceuticals.

In the following section, we discuss production and supply aspects of candidates that are currently in clinical practice ( $^{227}\text{Th}/^{223}\text{Ra}$ ,  $^{225}\text{Ac}$ ,  $^{211}\text{At}$ , and  $^{212}\text{Pb}/^{212}\text{Bi}$ ) and several promising candidates that are in preclinical evaluation ( $^{230}\text{U}/^{226}\text{Th}$  and  $^{149}\text{Tb}$ ). All nuclear decay data are taken from the NuDat library, version 2.8 (<https://www.nndc.bnl.gov/nudat2/>).

## CLINICALLY RELEVANT $\alpha$ -EMITTERS

### $^{227}\text{Th}/^{223}\text{Ra}$

Starting from  $^{227}\text{Ac}$  (half-life [ $t_{1/2}$ ], 21.77 y), 2 nuclides for TAT applications can be extracted:  $^{227}\text{Th}$  ( $t_{1/2}$ , 18.7 d) and  $^{223}\text{Ra}$  ( $t_{1/2}$ , 11.43 d) (Fig. 1). Despite their chemical differences, they are grouped because of their common starting material, similarly to  $^{225}\text{Ac}$  and  $^{213}\text{Bi}$  or  $^{224}\text{Ra}$  and  $^{212}\text{Pb}$ . Thus, before exploring the current use of  $^{227}\text{Th}$  and  $^{223}\text{Ra}$ , the production of  $^{227}\text{Ac}$  needs to be described.

$^{227}\text{Ac}$  is produced primarily via neutron irradiation of a  $^{226}\text{Ra}$  target in a nuclear reactor (12). Several limitations relevant to  $^{226}\text{Ra}$  ( $t_{1/2}$ , 1600 y) as a target material need to be considered: the target is highly radioactive, with the  $^{222}\text{Rn}$  ( $t_{1/2}$ , 3.82 d) daughter as a radioactive gas; further, limited quantities of  $^{226}\text{Ra}$  are currently available. For an efficient process, a high flux of thermal neutrons with minimum contribution of fast neutrons is preferable, as  $^{226}\text{Ra}$  does have a non-negligible fission cross-section for neutrons with energy above 1 MeV (13). Because of the increased focus of medical authorities on production quality, a specification of the target material may be required to ensure the quality of the  $^{227}\text{Ac}$  product.

After  $^{226}\text{Ra}$  target irradiation, purification of  $^{227}\text{Ac}$  from the target is the next step. This is accomplished by separation using liquid chromatography techniques, similar to the procedure for separation of  $^{229}\text{Th}$  from  $^{225}\text{Ac}$  and  $^{225}\text{Ra}$  (14). The separation process needs to remove all radium and all thorium, as both  $^{228}\text{Th}$  and  $^{229}\text{Th}$  will be present as by-products after irradiation, along with the remaining  $^{226}\text{Ra}$ . After  $^{227}\text{Ac}$  purification, characterization of the actinium is recommended, to have as much data on the starting material as possible. The  $^{227}\text{Ac}$  is typically kept in a dilute nitric solution but may be dried down to actinium nitrate if the material is to be shipped, because shipment of dry material is easier than shipment of a solution, based on the current International Air Transport Association regulations (15).

Alternative approaches include the recovery of  $^{227}\text{Ac}$  from legacy actinium-beryllium neutron sources (16) and the accelerator-based production of  $^{225}\text{Ac}$  using  $^{232}\text{Th}$  as a target generating small quantities of  $^{227}\text{Ac}$  as a by-product (17). There was virtually no production of  $^{227}\text{Ac}$  between the 1970s and the past decade. Thus, the availability of  $^{227}\text{Th}$  and  $^{223}\text{Ra}$  was very limited.

## NOTEWORTHY

- TAT shows significant potential in the clinic for cancer treatment.
- One of the main challenges limiting the wide application of TAT is the production and supply of suitable TAT radionuclides.
- For most TAT radionuclides, current demand significantly exceeds the available supply, but international efforts for increased production are under way.

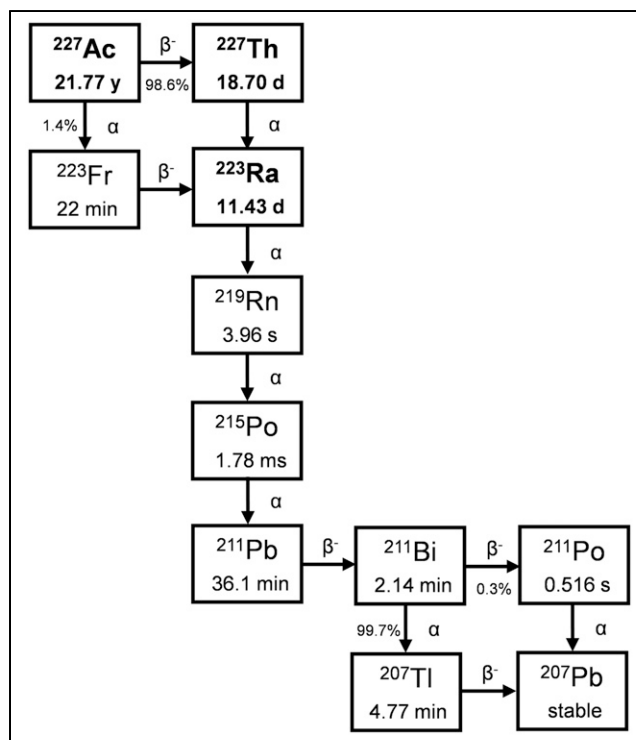


FIGURE 1. Decay scheme of  $^{227}\text{Ac}$ .

$^{227}\text{Th}$  is harvested from a generator containing  $^{227}\text{Ac}$ . Using separation columns, it is possible to separate thorium from actinium and radium, thus removing both the mother and the daughter nuclides. The purified thorium may be used on-site for immediate labeling or shipping, as thorium chloride, to the labeling site. If shipment or labeling is delayed, the purification step for removal of radium may be repeated to minimize the dose contribution from daughters.

$^{223}\text{Ra}$  is also harvested from a generator containing  $^{227}\text{Ac}$ . Using separation columns, radium can be separated from actinium and thorium, thus removing both mother nuclides. The purified radium is typically used on-site for drug formulation immediately or shipped as dry radium chloride.

Both  $^{223}\text{Ra}$  and  $^{227}\text{Th}$  are currently commercially available from Oak Ridge National Laboratory (ORNL) through the U.S. Isotope Distribution Office, and Pacific Northwest National Laboratory, Rosatom, and Bayer have access to  $^{223}\text{Ra}$  and  $^{227}\text{Th}$ .

Radium was considered a good candidate for TAT, but over the last few decades, no suitable chelator has been found. However,  $^{223}\text{Ra}$  in its ionic form is clinically used as Xofigo in the treatment of bone metastatic prostate cancer (18). This does not involve a chelator or a target-seeking moiety. Xofigo thus represents a special form of TAT pharmaceutical. The use and handling of Xofigo, which is currently approved in 53 countries, form the basis for any subsequent TAT pharmaceuticals, including those produced in the European Union, the United States, and Japan.

Because of their short half-lives, all daughters are in radioactive equilibrium with  $^{223}\text{Ra}$  at the time of injection. The shelf-life of a radiopharmaceutical will be governed by several parameters, including the activity of the mother nuclide, ingrowth of radioactive daughters, and degradation of the pharmaceutical due to

radiolysis of one or several components. In the case of Xofigo, the ingrowth of daughters is not a limiting factor, as the daughters are fully ingrown after some hours. Radiolysis of the pharmaceutical is a concern, particularly radiolysis of the citrate buffer. In addition, the lower specific radioactivity after several half-lives is also a concern. To determine a proper shelf-life, these aspects must be considered and studies must be conducted. In the case of Xofigo, a shelf-life of 28 d was therefore adopted. This shelf-life is unusually long for a radiopharmaceutical, partly because of the  $t_{1/2}$  of  $^{223}\text{Ra}$  and partly because of the low impact of radiolysis, and allows for global distribution independent of production site location. For research applications, the availability of  $^{227}\text{Ac}$ ,  $^{227}\text{Th}$ , and  $^{223}\text{Ra}$  is currently sufficient, but the overall supply for clinical or commercial use is less certain. No published data on the production capacity for the different suppliers are available.

### $^{225}\text{Ac}/^{213}\text{Bi}$

$^{225}\text{Ac}$  is one of the most promising TAT radionuclides, with a  $t_{1/2}$  of 9.92 d and a net emission of 4  $\alpha$ -particles in the decay chain. It can be used for TAT radiopharmaceuticals or as a source of  $^{213}\text{Bi}$  ( $t_{1/2}$ , 45.61 min), which also can be applied in TAT (19).

There are several production routes for  $^{225}\text{Ac}$  (20). The two most important are separation from the natural decay of  $^{229}\text{Th}$  obtained from waste stockpiles containing  $^{233}\text{U}$ , and irradiation of  $^{232}\text{Th}$  with high-energy protons ( $>70$  MeV) via reaction  $^{232}\text{Th}(p,x)^{225}\text{Ac}$ .

In addition, irradiation of  $^{226}\text{Ra}$  with lower-energy protons ( $<25$  MeV) via the reaction  $^{226}\text{Ra}(p,2n)^{225}\text{Ac}$  holds great promise for large quantities because of the high (710 mbarn) cross-section peak at 16.8 MeV (21). This reaction could be performed on many of the low-energy cyclotrons already in use for medical isotope production. However, irradiation of a highly radioactive target on these medical cyclotrons, and limited radium quantities, have rendered use of this approach infrequent. Another promising route is photonuclear production via  $^{226}\text{Ra}(\gamma,n)^{225}\text{Ra} \rightarrow ^{225}\text{Ac}$ , which can provide the clinically relevant supply of  $^{225}\text{Ac}$  (22).

### Production of $^{225}\text{Ac}$ from $^{229}\text{Th}$ Decay

$^{225}\text{Ac}$  is most frequently produced from  $^{229}\text{Th}$  generators.  $^{229}\text{Th}$  is the grandparent isotope of  $^{225}\text{Ac}$  in the decay series of  $^{233}\text{U}$  (Fig. 2) and has a  $t_{1/2}$  of 7,932 y (NuDat), later corrected to 7,917 y (23). As such,  $^{229}\text{Th}$  serves as an ideal radioisotope generator for a virtually perpetual supply of  $^{225}\text{Ac}$ . The primordial neptunium decay chain to which  $^{229}\text{Th}$  belongs is now extinct; therefore,  $^{229}\text{Th}$  that is suitable for use via separation technology is of limited availability. The most common source of  $^{229}\text{Th}$  is the decay of anthropogenic  $^{233}\text{U}$ . Because of safeguarding and nonproliferation efforts surrounding  $^{233}\text{U}$ , access to large quantities is limited and approximately only 12.9 GBq (350 mCi) of  $^{229}\text{Th}$  have been converted into functioning  $^{225}\text{Ac}$  generators to date; this restriction has limited the global annual production of  $^{225}\text{Ac}$  to approximately 63 GBq (1.7 Ci) (20).

By allowing the  $^{225}\text{Ra}$  ( $t_{1/2}$ , 14.9 d) and  $^{225}\text{Ac}$  ( $t_{1/2}$ , 9.92 d) progeny of  $^{229}\text{Th}$  to approach secular equilibrium (Supplemental Fig. 1; supplemental materials are available at <http://jnm.snmjournals.org>) over typically 30–90 d, the generator can be eluted to separate the shorter-lived daughters. After initial milking of the  $^{229}\text{Th}$  generator,  $^{225}\text{Ra}$  can be stored for further use as a parent–daughter generator of a reduced but still relevant quantity of  $^{225}\text{Ac}$ . The frequency of  $^{229}\text{Th}$  and  $^{225}\text{Ra}$  generator elution is

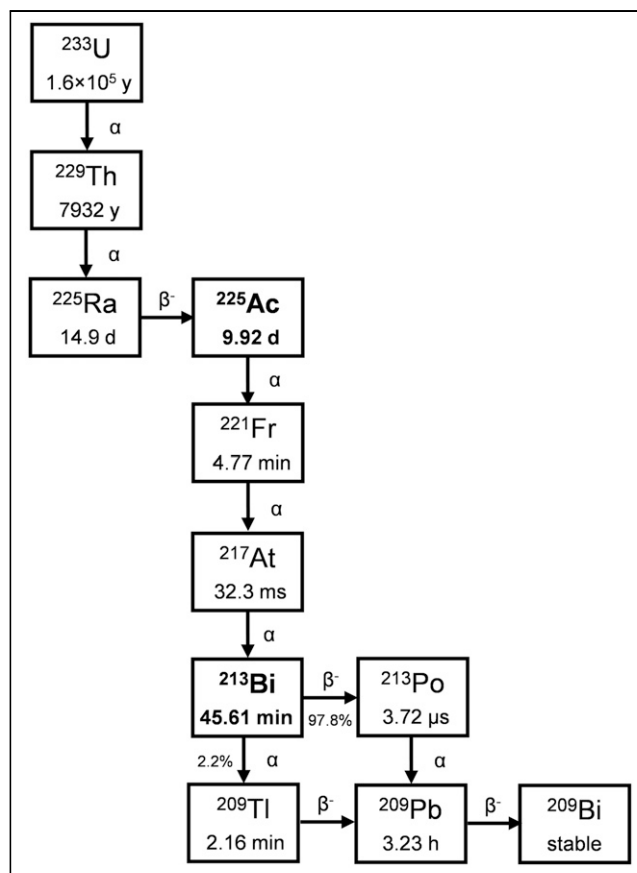


FIGURE 2. Decay scheme of  $^{233}\text{U}$ .

often determined with considerations for batch size requirements, operational cost, and generator size. Generators with larger amounts of  $^{229}\text{Th}$  can produce suitable  $^{225}\text{Ac}$  batch sizes with higher frequency. After elution, selective isolation and careful quality control are performed to prepare  $^{225}\text{Ac}$  suitable for incorporation into radiopharmaceuticals.

There are multiple  $^{229}\text{Th}$  generators in operation, capable of producing  $^{225}\text{Ac}$  in quantities relevant for preclinical and limited clinical use. The Directorate for Nuclear Safety and Security of the Joint Research Centre in Karlsruhe, Germany (Supplemental Fig. 2), possesses about 215 mg of  $^{229}\text{Th}$  (24), and the Leypunsky Institute for Physics and Power Engineer in the Russian Federation (Supplemental Fig. 3) (25) and ORNL in the United States (26) each possesses 700 mg of  $^{229}\text{Th}$ . An additional source recently became available at Canadian Nuclear Laboratories (Supplemental Fig. 4) (27), with 50 mg of  $^{229}\text{Th}$ .

All generators rely on an anion exchange mechanism for separation of  $^{229}\text{Th}$  from  $^{225}\text{Ra}$  and  $^{225}\text{Ac}$ . Cation exchange or extraction chromatography is used for  $^{225}\text{Ac}$  separation from  $^{225}\text{Ra}$ , whereas additional anion exchange processing provides purification from residual thorium (28). These methods produce  $^{225}\text{Ac}$  with suitable attributes for preclinical and clinical applications (7). Molar-specific activity and stable metal content can differ for various  $^{225}\text{Ac}$  sources.  $^{233}\text{U}$  stockpiles are currently being processed at ORNL through a public–private partnership that is expected to yield about 45,000 mg of  $^{229}\text{Th}$  (Supplemental Table 1) (29). This material also contains  $^{228}\text{Th}$  in sufficient quantities (30) to require

exposure shielding due to the presence of  $^{208}\text{Tl}$ , complicating generator development and deployment. Work on the generator for this material is ongoing and will leverage the techniques and methods used in the existing generators (31).

### Production of $^{225}\text{Ac}$ via Proton Irradiation of $^{232}\text{Th}$

$^{225}\text{Ac}$  has been produced by the spallation reaction  $^{232}\text{Th}(p,x)^{225}\text{Ac}$  on thorium targets with proton energies ranging from 100 to 1,400 MeV at beam currents of as high as 250  $\mu\text{A}$ . This method of production is presently under development as part of the U.S. Department of Energy's (DOE's) Tri-Lab Effort, involving Brookhaven National Laboratory and Los Alamos National Laboratory for target irradiations and ORNL for subsequent radiochemical processing and dispensing of irradiated targets. The current focus of the U.S. DOE Tri-Lab Effort is to bring colocated processing facilities online at Brookhaven National Laboratory and Los Alamos National Laboratory in the time frame of 6 mo to 5 y, with the goal of greater than monthly production of curie-scale batches. The U.S. DOE Tri-Lab Effort has established processing under current good-manufacturing-practice conditions, with operations captured under a drug master file (32,33).

In addition, efforts are under way to develop spallation production capabilities in Canada using a diverse set of irradiation capabilities at the Tri-University Meson Facility (TRIUMF) (up to 500-MeV proton beams). Producing  $^{225}\text{Ac}$  at higher proton energy results in a higher fraction of  $^{225}\text{Ac}$  versus  $^{227}\text{Ac}$  (Supplemental Fig. 5). This is currently being pursued with TRIUMF's 500-MeV cyclotron. About 200-MBq (5.4 mCi) quantities of  $^{225}\text{Ac}$  are produced with a proton beam of about 480 MeV on target. In addition, significant amounts of  $^{225}\text{Ra}$  are also produced, which can be separated and used as a generator isotope for isotopically pure  $^{225}\text{Ac}$  (Supplemental Fig. 5). In recent 25 mA·h irradiations of approximately 8-g targets of  $^{232}\text{Th}$ , isolation of the radium fraction provided sufficient  $^{225}\text{Ra}$  to yield about 18 MBq ( $\sim 0.5$  mCi) of  $^{225}\text{Ac}$ , with no detectable  $^{227}\text{Ac}$  (Supplemental Fig. 6 shows the separation, and Supplemental Fig. 7 shows an example of a  $\gamma$ -spectrum (34,35)).

Furthermore, work is under way at the Institute of Nuclear Research, Russia, and at NorthStar Medical Technologies, where teams are developing spallation production targets and new process technologies (36–38).

In thorium spallation,  $^{227}\text{Ac}$  ( $t_{1/2}$ , 21.77 y) is coproduced with yields similar to those for  $^{225}\text{Ac}$ , leading to concerns about facility licensing and about the path forward for associated waste streams. Overall,  $^{227}\text{Ac}$  activity represents approximately 1%–2% of the overall sample activity and has not been demonstrated to impact labeling efficiency with DOTA, the gold standard for radiolabeling, or to result in toxicity concerns (39). The concern related to  $^{227}\text{Ac}$  content can be avoided when producing  $^{213}\text{Bi}$  from a generator, which retains all actinium isotopes. This issue can also be addressed by isolating radium isotopes and further extracting isotopically pure  $^{225}\text{Ac}$  (40). Additionally, researchers at TRIUMF have demonstrated online generation of isotopically pure beams of  $^{225}\text{Ac}$  using a resonant laser ionization method (41), and Conseil Européen pour la Recherche Nucléaire (CERN) has demonstrated separation of pure beams from different thick targets of either  $^{225}\text{RaF}^+$  or  $^{225}\text{Ac}^+$ , including molecular ion formation or resonant laser ionization (11). The supply of  $^{225}\text{Ra}$  or  $^{225}\text{Ac}$  from CERN-MEDICIS (Medical Isotopes Collected from Isolde) or Institute for Transuranium Elements, Karlsruhe, will become available for researchers through a newly approved coordinated European hub,

PRISMAP (Production of High Purity Isotopes by Mass Separation for Medical Application). The European medical isotope program kicked off in 2021 (42).

Currently, most  $^{225}\text{Ac}$  is used in the form of  $^{225}\text{Ac}$ -labeled radiopharmaceuticals (43), both for preclinical development and for clinical studies mainly focusing on the treatment of prostate cancer, neuroendocrine tumors, and gliomas. Although the application of  $^{213}\text{Bi}$ , generated from  $^{225}\text{Ac}/^{213}\text{Bi}$  generators, has also demonstrated significant clinical benefit (44), the limited availability and high cost of high-activity generators are presently hampering further studies.

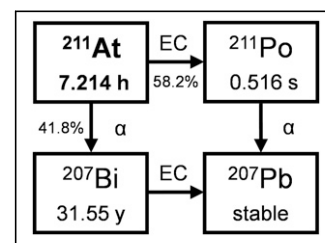


FIGURE 3. Decay scheme of  $^{211}\text{At}$ .

### $^{211}\text{At}$

$^{211}\text{At}$  (Fig. 3) is the  $\alpha$ -emitting radionuclide that is perhaps easiest to produce. However, its availability has been limited because there are few accelerators in the world that produce an  $\alpha$ -beam with the optimal energy range (28–29 MeV) and beam current (10  $\mu\text{A}$  or higher) to produce adequate quantities for research and clinical applications (5,45,46). Additionally, the relatively short  $t_{1/2}$  of  $^{211}\text{At}$  (7.21 h) causes distribution problems. Here, the supply model is a network, as, for instance, established by the DOE isotope program.

The most common method of production uses the  $^{209}\text{Bi}(\alpha,n)^{211}\text{At}$  reaction, in which a metallic bismuth target is bombarded with  $\alpha$ -particles (Fig. 4). Inexpensive naturally abundant monoisotopic bismuth (available at 99.999%) can be used directly for target preparation. In general, the bismuth metal is melted onto or is deposited from a vapor onto an aluminum or copper backing. Bismuth metal, although inexpensive, is a poor thermal conductor and has a low melting point (272°C). Thus, effective cooling methods to prevent the target from melting during irradiation are required (47). Thick targets (80  $\mu\text{m}$ ) are desired for production and to keep the beam from hitting the target backing, but thinner targets allow for the most efficient cooling. Alternate bismuth target materials with higher melting points, such as  $\text{Bi}_2\text{O}_3$ , can be used in irradiation, but thus far none have proven to be superior to bismuth metal in the production of  $^{211}\text{At}$ .

The incident energy of the  $\alpha$ -beam in bismuth irradiations is important for optimizing  $^{211}\text{At}$  production rates and for

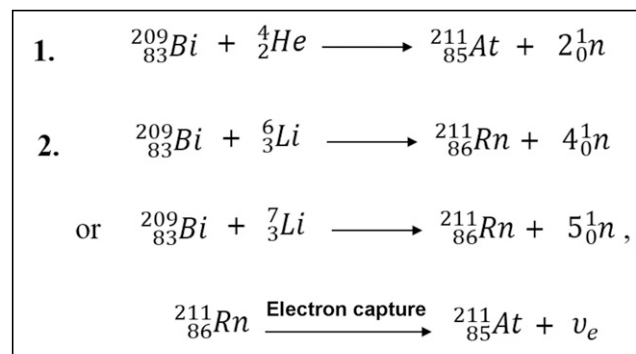


FIGURE 4. Direct and indirect production routes of  $^{211}\text{At}$ .  $^{211}\text{Rn}$  can also be produced by spallation of actinide targets (uranium or thorium) induced by high-energy protons (reaction not shown).

minimizing production of an unwanted radionuclide,  $^{210}\text{At}$ . Production of  $^{210}\text{At}$  ( $t_{1/2}$ , 8.1 h) is problematic as it decays to a long- $t_{1/2}$   $\alpha$ -emitter,  $^{210}\text{Po}$  ( $t_{1/2}$ , 138.38 d). Although  $^{210}\text{Po}$  is found in nature, it has high human toxicity. Generally, a 28-MeV  $\alpha$ -beam has been used to preclude  $^{210}\text{At}$  production. In an optimization study at 29 MeV, no quantities of  $^{210}\text{At}$  were detected (48). At 29 MeV, the production rate of  $^{211}\text{At}$  was increased by about 15% over that of a 28-MeV irradiation. Production of  $^{211}\text{At}$  can be significantly increased by increasing the accelerator beam current, but unfortunately, the feasible beam current is inherent in the design of the accelerator. However, a higher beam current can be obtained by irradiating an internal target as the beam current is lost during extraction to external beamlines (49).

Isolation of pure  $^{211}\text{At}$  from irradiated bismuth targets is also relatively simple compared with other  $\alpha$ -emitters, as there are no other radionuclides produced under optimal irradiation conditions. The most common and perhaps simplest method for isolation of  $^{211}\text{At}$  is high-temperature (650°C–700°C) dry distillation. However, there can be radiation safety concerns with volatilized  $^{211}\text{At}$ . Therefore, alternative wet chemistry isolation methods are being developed (50,51). To simplify the isolation of  $^{211}\text{At}$ , methods for automation of dry distillation and wet chemistry approaches are being developed (52).

Current quantities of  $^{211}\text{At}$  are inadequate for widespread clinical use. In fact, only Duke University and the University of Washington in the United States, and Copenhagen University Hospital in Denmark, have produced  $^{211}\text{At}$  for clinical trials.

The U.S. DOE Office of Isotope Research and Development and Production provides funding to improve the availability of  $^{211}\text{At}$  in the United States. It is also creating a university network for  $^{211}\text{At}$  production in different regions of the United States for shipment to users through the DOE National Isotope Development Center. Japan has 5 sites producing  $^{211}\text{At}$  by  $\alpha$ -beam irradiation for use at 13 sites. In support of research, the European Union has recently initiated a cost action (CA19114) that involves networking of  $^{211}\text{At}$  production centers among several European countries.

Although production of  $^{211}\text{At}$  is currently limited, accelerators with medium-energy  $\alpha$ -beams can be added at a significantly lower cost than high-energy accelerators or nuclear reactors required for the production of other  $\alpha$ -emitters. Accelerator technology innovations could provide much higher  $\alpha$ -beam currents than do current systems. Although new target technology will be required with higher  $\alpha$ -beam currents to circumvent target melting, ultimately much larger quantities of  $^{211}\text{At}$  could be produced.

An alternative, early-stage, research approach for  $^{211}\text{At}$  production involves irradiation of bismuth metal with lithium ions to produce  $^{211}\text{Rn}$  (Fig. 4) for a  $^{211}\text{Rn}/^{211}\text{At}$  generator. Since  $^{211}\text{Rn}$  has a  $t_{1/2}$  of 14.6 h, its decay during transit might provide a more effective distribution of  $^{211}\text{At}$ . Radon is classified as a gaseous element and may not be suitable for chemical operations. However, since it has a high affinity for nonpolar organic solvents, it is possible to make a generator by applying the solvent extraction method (53). Such a generator system has been demonstrated in which gaseous  $^{211}\text{Rn}$  was isolated and retained in liquid alkane hydrocarbon (dodecane), and  $^{211}\text{At}$  generated from the  $^{211}\text{Rn}$  source was extracted in an aqueous solution (2N NaOH) (54).

#### $^{212}\text{Pb}/^{212}\text{Bi}$

Both  $^{212}\text{Bi}$  ( $t_{1/2}$ , 60.55 min) and  $^{212}\text{Pb}$  ( $t_{1/2}$ , 10.64 h) are part of the  $^{232}\text{Th}$  ( $t_{1/2}$ ,  $1.4 \times 10^{10}$  y) and  $^{232}\text{U}$  ( $t_{1/2}$ , 68.9 y) decay chain, with  $^{212}\text{Bi}$  being the decay daughter of  $^{212}\text{Pb}$  (Fig. 5).  $^{212}\text{Pb}$  emits

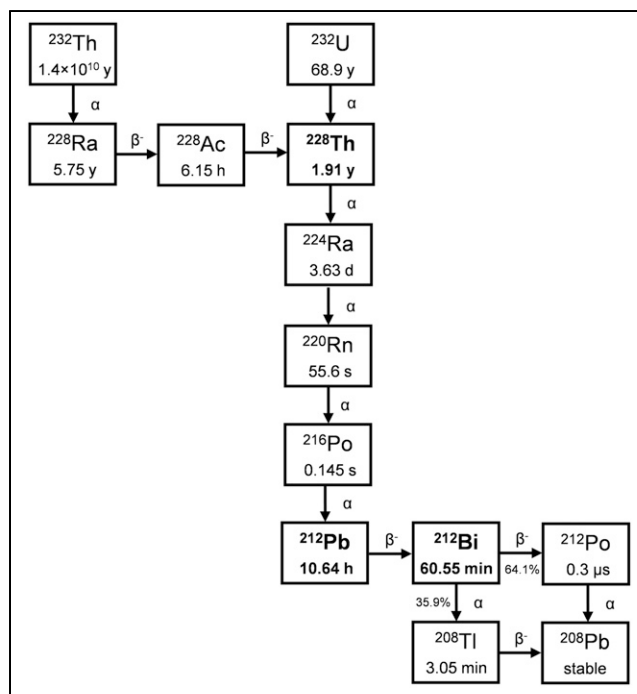


FIGURE 5. Decay scheme of  $^{232}\text{Th}$  and  $^{232}\text{U}$ .

2  $\beta^-$ -particles and 1  $\alpha$ -particle through its decay to stable  $^{208}\text{Pb}$ , whereas  $^{212}\text{Bi}$  emits 1  $\beta^-$ -particle and 1  $\alpha$ -particle, which can be used for targeted radionuclide therapy. Either radionuclide is commonly isolated from  $^{228}\text{Th}$  sources, which is a decay daughter of both  $^{232}\text{Th}$  and  $^{232}\text{U}$ .

A major drawback to using  $^{212}\text{Bi}$  clinically is the emission of a relatively intense and very high energy  $\gamma$ -ray (2.6 MeV of 36% intensity per decay of  $^{212}\text{Bi}$ ) via its daughter  $^{208}\text{Tl}$ . This also creates an obstacle to handling  $^{228}\text{Th}$  sources, leading to stability issues due to radiolytic damage of generator systems, and mandates significant shielding for operators (55). These issues have been noted for the newly available  $^{229}\text{Th}$ , which contains  $^{228}\text{Th}$  in relatively high quantities (as described in the “ $^{225}\text{Ac}/^{213}\text{Bi}$ ” section).

Using  $^{212}\text{Pb}$  as an in vivo generator, instead of  $^{212}\text{Bi}$  directly, in radiopharmaceutical development reduces the amount needed for therapy 10-fold and facilitates radiopharmaceutical production, formulation, and administration given its longer  $t_{1/2}$ . However, the issue of daughter recoil after  $\beta$ -decay and subsequent retention of progeny needs to be taken into consideration (56), similarly to the other  $\alpha$ -emitters discussed.

$^{212}\text{Pb}$ , and thus  $^{212}\text{Bi}$ , are isolated from  $^{228}\text{Th}$  or  $^{224}\text{Ra}$  generators, both of which are natural decay products of  $^{232}\text{Th}$ .  $^{228}\text{Th}$  can be obtained through isolation of  $^{228}\text{Ra}$  at annual intervals from  $^{232}\text{Th}$  or by isolation from anthropologic sources via  $^{232}\text{U}$  stockpiles (a portion of which has been transferred by the Department of Defense to AlphaMed, Inc., from stocks at ORNL, or by the double-neutron capture and successive  $\beta^-$  decay of  $^{226}\text{Ra}$  (57)).

Isolation from natural  $^{228}\text{Ra}$  remains difficult given the need to process tons of aged  $^{232}\text{Th}$  to obtain useable amounts. Each ton of more than 35-y-old  $^{232}\text{Th}$  can yield approximately 3.7 GBq (100 mCi) of  $^{228}\text{Ra}$ . The company OranoMed extracts  $^{212}\text{Pb}$  from natural thorium salt. Subsequent separation, purification, and concentration of elements decaying from  $^{232}\text{Th}$  provide worldwide shipment of  $^{212}\text{Pb}$  generators.

$^{228}\text{Th}$  can be produced from successive neutron capture and  $\beta^-$  decay of  $^{226}\text{Ra}$ . In the past, this production was proven to be feasible, but further process development is needed to determine production yields and cost.

Around 555-MBq (15 mCi)  $^{224}\text{Ra}/^{212}\text{Pb}/^{212}\text{Bi}$  generators are available through the U.S. DOE Isotope Program (via ORNL) (58). The current generator is sufficient only for preclinical development (not clinical use), as radiolytic damage limits the scale-up (59). Briefly,  $^{224}\text{Ra}$  ( $t_{1/2}$ , 3.63 d) is separated from immobilized  $^{228}\text{Th}$  adsorbed onto an organic cation exchange resin (highly cross-linked MP-50,  $\sim 300\ \mu\text{L}$  in volume). A  $^{212}\text{Pb}$  and  $^{212}\text{Bi}$  mixture is eluted with a few milliliters of 2 M HCl or 0.5 M HI with approximately 70% yield and parent breakthrough of  $10^{-6}$ . It is also possible to elute  $^{212}\text{Bi}$  (free from  $^{212}\text{Pb}$ ) selectively with 0.5 M HCl or 0.15 M HI. The  $^{224}\text{Ra}/^{212}\text{Pb}/^{212}\text{Bi}$  generator has a shelf-life of about 2 wk.

Weström et al. (60) prepared a  $^{228}\text{Th}/^{224}\text{Ra}$  generator based on thorium purchased from Eckert and Ziegler. In this process,  $^{228}\text{Th}$  was immobilized on a DIPEX (Eichrom) actinide resin by mixing  $^{228}\text{Th}$  in 0.1 M  $\text{HNO}_3$  with a portion of the actinide resin and, after a few hours, loading onto a column containing a small portion of inactive actinide resin to avoid breakthrough.  $^{224}\text{Ra}$  could be eluted regularly from the generator column with 1 M HCl.

McNeil et al. (61) reported the preparation of a novel  $^{228}\text{Th}/^{212}\text{Pb}$  generator using  $^{228}\text{Th}$  produced as a by-product of  $^{232}\text{Th}$  spallation with 500-MeV protons at TRIUMF. The bulk thorium (8 g) (coprecipitated with  $^{228}\text{Th}$ ) was purified via anion exchange resin.  $^{228}\text{Th}$  did not absorb to the column and was found in load and wash fractions, which were collected, evaporated to dryness, and redissolved in 1 M  $\text{HNO}_3$  to produce the generator stock solution. The  $^{212}\text{Pb}$  was separated from  $^{228}\text{Th}$  by passing the generator stock solution through an 80-mg lead resin (Eichrom).

## Research Candidates

There are several hundred  $\alpha$ -emitters in the chart of radionuclides; however, most are not suitable for TAT because of their  $t_{1/2}$  or difficulties with the production and formulation of radiopharmaceuticals. However, with emerging alternative production routes and advancement in chelation systems, several additional candidates are of interest for TAT.

### $^{230}\text{U}/^{226}\text{Th}$

$^{230}\text{U}$  ( $t_{1/2}$ , 20.8 d) can be used for TAT directly or as a generator source of shorter-lived  $^{226}\text{Th}$  ( $t_{1/2}$ , 30.57 min). One potential advantage over  $^{225}\text{Ac}/^{213}\text{Bi}$  is that  $^{230}\text{U}/^{226}\text{Th}$  has multiple  $\alpha$ -decays with very short lived daughters (seconds), which potentially may prevent significant delocalization of daughters after the decay from the targeting site (Fig. 6). One of the main challenges in using  $^{230}\text{U}$  in the same way as  $^{225}\text{Ac}$  for direct labeling of biomolecules (e.g., antibodies or peptides) is the still relatively undeveloped chelation of uranium for radiopharmaceutical application. In addition, for the  $^{230}\text{U}/^{226}\text{Th}$  generator, the shorter  $t_{1/2}$  of  $^{226}\text{Th}$  may represent challenges in term of logistic and radiopharmaceutical synthesis when used in the same way as  $^{213}\text{Bi}$ .  $^{230}\text{U}$  can be produced either directly by proton or deuteron irradiation of  $^{231}\text{Pa}$  ( $t_{1/2}$ ,  $3.276 \times 10^4\ \text{y}$ ) (62,63) or by decay of  $^{230}\text{Pa}$  ( $t_{1/2}$ , 17.4 d), which can be produced by spallation of  $^{232}\text{Th}$  (64–66). For the first route, the production rate is 0.25 MBq (6.7  $\mu\text{Ci}$ )/ $\mu\text{Ah}$  for 25 MeV of energy (65), and limiting factors are the handling and availability of the target material. For the second route, the main limitation is that only 7.8% of produced  $^{230}\text{Pa}$  decays to  $^{230}\text{U}$ , significantly

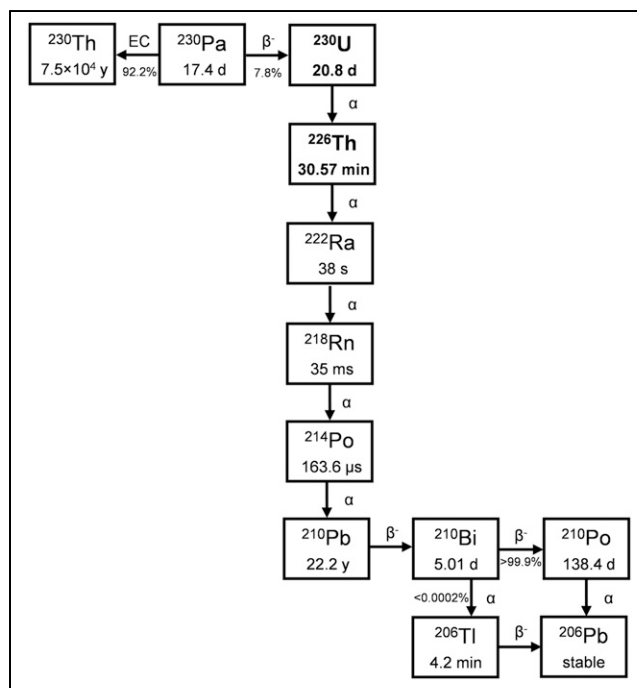


FIGURE 6. Decay scheme of  $^{230}\text{Pa}$ .

decreasing the final yield.  $^{230}\text{Pa}$  can also be produced as a by-product during proton spallation of  $^{232}\text{Th}$  and coextracted along with other medical radionuclides (67), such as  $^{225}\text{Ac}$ . Radiochemical processing is required for the protactinium, uranium, and thorium, as is already well known and can be achieved by a combination of ion-exchange and solid-phase extraction chromatography. On the basis of the similar required clinical quantities of  $^{225}\text{Ac}$ ,  $^{230}\text{U}$  can be produced (gigabecquerels [tens of millicuries]) via both routes; however, the currently low demand and absence of a suitable chelation system for uranium limit application of this attractive radionuclide.

### $^{149}\text{Tb}$

$^{149}\text{Tb}$  ( $t_{1/2}$ , 4.118 h) is the lightest  $\alpha$ -emitter if one excludes those with the extremely short ( $^{108}\text{Te}$ ) or long ( $^{146}\text{Sm}$ )  $t_{1/2}$ .  $^{149}\text{Tb}$  was recognized long ago as an  $\alpha$ -emitting radionuclide with potentially interesting properties (Fig. 7), as it was found to be capable of killing single cancer cells in vitro and is part of a terbium therapeutic quadruplet covering the different nuclear medicine modalities in therapy and diagnostics (68–70). Cyclotron irradiation of gadolinium targets with high-energy protons (70–200 MeV) and spallation reactions of protons at high energy ( $>1\ \text{GeV}$ ) on thick tantalum targets are the most favorable production routes. In all cases, mass separation is required to suppress coproduced terbium

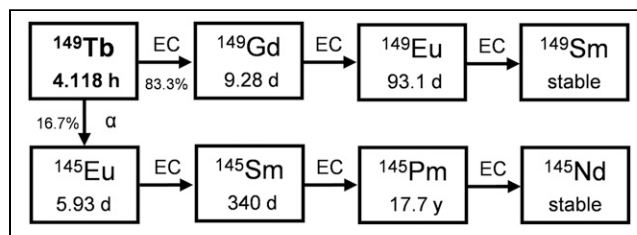


FIGURE 7. Decay scheme of  $^{149}\text{Tb}$ .



radionuclides, with demonstrated efficiencies of 12% at CERN-MEDICIS and of 50% with stable terbium tracers at the LARISSA (Laser Resonance Ionization Spectroscopy for Selective Applications) isotope separator at the University of Mainz (Germany) and for the  $^{149}\text{Tb}$  production route followed at ISOLDE (Isotope Mass Separator On-Line). Daily cycles of 500-MBq batches are expected from 2021 onward at CERN-MEDICIS to produce no-carrier-added  $^{149}\text{Tb}$  radionuclide batches.

The relatively short  $t_{1/2}$  of  $^{149}\text{Tb}$  implies distribution networks mimicking those of many diagnostic radiopharmaceuticals. To support  $^{149}\text{Tb}$  distribution, transportation limits have been updated accordingly in 2018 and are no longer a limiting factor for the dispatch of relevant activities for clinical applications (71).

Although these different  $\alpha$ -emitters have been made available to researchers under different access modalities, a new consortium has been established, funded by the European Union's Horizon 2020 research and innovation program. PRISMAP comprises important nuclear reactors, accelerators, and isotope mass separation centers. It aims at providing different radionuclides for medical researchers through a single hub, a single web platform (42). A call for projects, selection by a user panel, and determination of important nuclear data for proper standardization are fully included in the project's implementation. PRISMAP started on May 1, 2021.

## DISCUSSION

The current supply of most TAT radionuclides is insufficient for preclinical and clinical evaluation. Only very few research groups have reliable access to TAT radionuclides, because of either high costs or long wait times. Therefore, the supply of  $\alpha$ -emitting radionuclides for TAT is a pressing issue that needs to be addressed urgently. The current review intends to stimulate discussion and provide useful information on the selection and handling of TAT radionuclides for scientists and clinicians who would like to develop TAT programs. Table 1 summarizes the nuclear

properties, current availability on a clinical scale, and the potential for a future increase in production.

Several strategies can potentially solve the supply shortage of TAT radionuclides: development of alternative production strategies by exploiting existing infrastructure or developing novel approaches; improvement of targeting and radiochemical separation strategies enabling scaling up of the production of TAT radionuclides; achieving a fieldwide consensus on the appropriate radionuclidic and radiochemical purity for preclinical and clinical applications; development of a broad portfolio of different chelation and delivery systems to mitigate the physical and chemical limitations of  $\alpha$ -emitters; and definition of dosimetry and quality standards for clinical applications.

## CONCLUSION

The supply and production of  $\alpha$ -emitters are of critical importance when selecting the most suitable candidates for preclinical and, in particular, clinical TAT. The availability of  $\alpha$ -emitters has slowed the successful development of radiopharmaceuticals for TAT. Nevertheless, several landmark developments, including the success of Xofigo and  $^{225}\text{Ac}$ -labeled prostate-specific membrane antigen ligands, demonstrated the great potential of TAT. The strong academic and industry interest further stimulated by these success stories is expected to significantly improve radiopharmaceutical supply soon.

## DISCLOSURE

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**TABLE 1**  
Nuclear Properties of Discussed TAT Radionuclides and Their Current Production Route\*

| Isotope                           | Half-life         | Current production routes   | Potential to increase production  |
|-----------------------------------|-------------------|---|---|
| $^{227}\text{Th}/^{223}\text{Ra}$ | 18.70 d/11.43 d   | Decay of $^{227}\text{Ac}$ (generator $^{227}\text{Ac}/^{227}\text{Th}/^{223}\text{Ra}$ )   | Produce $^{227}\text{Ac}$ via neutron irradiation of $^{226}\text{Ra}$  |
| $^{225}\text{Ac}/^{213}\text{Bi}$ | 9.92 d/45.61 min  | Decay of $^{229}\text{Th}$ (generator $^{229}\text{Th}/^{225}\text{Ra}/^{225}\text{Ac}$ ); $^{232}\text{Th}(\text{p},\text{x})^{225}\text{Ac}$ ( $^{227}\text{Ac}$ contamination or $^{225}\text{Ra}/^{225}\text{Ac}$ generator)  | Provide additional stock of $^{229}\text{Th}$ ; scale up spallation on $^{232}\text{Th}$ production; $^{226}\text{Ra}(\text{p},2\text{n})^{225}\text{Ac}$ ; $^{226}\text{Ra}(\gamma,\text{n})^{225}\text{Ra} \rightarrow ^{225}\text{Ac}$ |
| $^{211}\text{At}$                 | 7.21 h            | $^{209}\text{Bi}(\alpha,2\text{n})^{211}\text{At}$  | Explore production at existing and upcoming facilities with $\alpha$ -beam; $^{211}\text{Rn}/^{211}\text{At}$ generator route   |
| $^{212}\text{Pb}/^{212}\text{Bi}$ | 10.64 h/60.55 min | Decay of $^{228}\text{Th}$ (generator $^{228}\text{Th}/^{224}\text{Ra}/^{212}\text{Pb}/^{212}\text{Bi}$ )   | Increase production of $^{228}\text{Th}$ (e.g., by-product of $^{227}\text{Ac}$ production and $^{232}\text{Th}$ spallation)  |
| $^{230}\text{U}/^{226}\text{Th}$  | 20.8 d/30.57 min  | $^{232}\text{Th}(\text{p},3\text{n})^{230}\text{Pa} \rightarrow ^{230}\text{U}/^{226}\text{Th}$ ; $^{231}\text{Pa}(\text{p},2\text{n})^{230}\text{U}/^{226}\text{Th}$ ; $^{232}\text{Th}(\text{p},\text{xn})^{230}\text{Pa} \rightarrow ^{230}\text{U}/^{226}\text{Th}$ | Develop scale-up production for p,3n route; extraction as by-product of $^{232}\text{Th}$ spallation  |
| $^{149}\text{Tb}$                 | 4.12 h            | $^{\text{nat}}\text{Ta}(\text{p},\text{x})^{149}\text{Tb}$ (mass separation)  | Produce regularly at CERN MEDICIS with PRISMAP initiative; engage other ISOL facilities   |

\*None have sufficient availability for routine clinical application.



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