

Innovative Facility for Isotope GENeration with Efficient Ion Accelerator

T4.2 Identify best radionuclides for production with LINAC

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T4.2 Identify best Isotopes for production with LINAC (M5-M42) [Leader: UL]

- 1. Analyising of precise nuclear data
- 2. for the selection of suitable radionuclides (target materials);
- 3. Investigating chemical forms and separation methods (high yields, activity scale, radiation stability), (+technical concept of the target materials)
- 4. Considering radiation safety and material recovery



T4.2 Identify best Isotopes for production with LINAC (M5-M42) [Leader: UL]

FOCUS:

- A. Radiolanthanides as main candidates? Others?
- B. Aim to support preclinical research and clinical uptake (the relevance and robustness of performed preclinical experiments) to <u>simplify their supply and utilization in the clinics</u>.
- C. Expand reactor-based production (not cyclotrons LINAC)
- D. Identify both established and novel radionuclides



T4.2 Identify best Isotopes for production with LINAC

Start Date:		M5	Task Leader:		UL		
End Date:		M15-42-48	Task Contribut	tors:	UL, ?		
Del.	Deliverab	le Title			Lead Partner	Diss. Level	Due On
D4.2	Report for the identification of the best candidateULSEN - Sensitive15 (42)isotopes for each of the three application areas:theranostics, therapy and diagnosticsthe sensitivethe sensitivethe sensitive						
Мх	Mileston	e Title		Lead Par	tner N	Aean of verification	Due On
4	Pilot pre-	clinical studies final	zation	6 - GNP		D4.3	48



- 1. Stepwise Approach: Data Collection Nuclear data (LINAC: cross-sections, halflife, emissions)
- 2. Feasibility Assessment LINAC energy & target compatibility
- 3. Radionuclide Prioritization Focus on:

Therapeutic radionuclides (e.g. Lu-177, Tb-161 etc.?)

Diagnostic emitters

Theranostic pairs (main focus in UL partners)

- 4. Process Optimization Target design, chemical form, separation
- 5. Validation & Selection Recommend radionuclides for pilot testing



Why LINAC?*

Reactor production is declining, needs are increasing LINACs are compact, tunable, and eco-friendly Enable local production in hospital or R&D settings Address the supply gap for *medium/long-lived* radionuclides

New radionuclides may open doors for next-gen radiopharmaceuticals

*Inspired by the NUSANO LINAC model (starting 2025)



Targeted Radionuclides to Discuss:

Lu-177: well-established therapeutic, growing demand Mo-99/Tc-99m: traditionally reactor-based generation Tb-161: longer-lived promising alternative for therapy (compared vs. cyclotron generated ones) Y-86/Y-90, Sc-44/Sc-47: theranostic options Potential new radionuclides (to be evaluated based on LINAC feasibility) ➡ Focus on beta emitters, theranostic pairs, longer-lived radionuclides

(compared vs. cyclotron generated ones)

Limit selection to 1–2 new isotopes for feasibility



• Task 4.2 Timeline: M5 – M42

Phase	Months	Description
Survey & selection	M5–M12	Literature, cross-sections, database
Feasibility study	M13–M24	LINAC compatibility
Separation strategy	M18–M30	Chemistry, recovery
Final selection	M30–M36	Candidate isotopes for pilot
Reporting	M36–M42	Deliverables, protocol input to WP4

Resources Involved:UL, WP3 coordination, inputs to/from T4.1–T4.5



- Collaboration is key:
- WP3: LINAC design and target irradiation feasibility
- **T4.1**: Legal and regulatory harmonization laboratory safety
- T4.3–T4.5: Radiochemistry, preclinical studies, clinical translation
- WP5: Strategic investment (scaling up production)
- Deliverable contributions:
- Protocols for Lu-177 and Mo-99
- Input for new radionuclides validation



• From the **Slovenian team's standpoint**:

LINAC does not replace cyclotron
LINAC can expand reactor-based production of new radionuclides
Aim to introduce new radionuclides – potential for theranostic use
Must consider both known radionuclides (Lu-177, Mo-99) and emerging ones

• **©** Goal: Practical, feasible set of radionuclides for the lab (WP4)





- 1. Coordinate with WP3 to get technical LINAC parameters:
 - technical parameters that <u>define LINAC-produced</u> radionuclides
 - 2. Define technical radionuclide shortlist focus on availability & impact
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- **3.** Cross-check with known medical demand (interviews?)
- 4. Contribute to planning pilot production protocols (T4.3–T4.5)
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- 5. Align with regulatory & infrastructural planning (T4.1, WP5)