

2026 JULY 19-22
VANCOUVER, BC
JOINT AAPM/COMP MEETING

Case Report of a Clinically Significant Lu-177 Extravasation: Rapid Clearance Kinetics and Dosimetry Using Quantitative SPECT/CT

Cancer
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ABSTRACT

Extravasation of Lu-177 radiopharmaceuticals can retain a large fraction of administered activity at the injection site, producing high local doses and reducing delivery to target tumors.

In this case of early extravasation during 200 mCi Lu-177 DOTATATE therapy, the infusion was stopped, IV access re-established, and clearance aided by local hyperthermia with arm elevation. Infiltrated activity was quantified with both a handheld CZT detector and quantitative SPECT/CT, yielding consistent estimates of 62–67 mCi within a 1070 mL soft-tissue volume.

Serial SPECT/CT imaging over 4.57 hours showed rapid clearance (effective half-life 3.66 hours) and a mean self-absorbed dose of 1.035 Gy to the compartment. The two low-dose CT scans contributed an additional 15 mSv effective dose, while the short clearance half-life avoided any change to standard patient release criteria.

This case illustrates a practical clinical workflow for timely extravasation assessment and dosimetry, demonstrating that prompt intervention and rapid biological clearance can limit tissue exposure even after substantial infiltration

INTRODUCTION

Extravasation during radiopharmaceutical therapy can retain a substantial fraction of administered activity at the injection site, creating high local absorbed doses and potential tissue injury while reducing delivery to target tumors.

For Lu-177 therapies, timely quantitative assessment of infiltrated activity, clearance, and extravasation dosimetry is essential to support patient-management decisions by quantifying the absorbed dose at the affected site, assessing the risk of tissue damage, supporting documentation, and guiding timely interventions to mitigate harm [1].



Figure 2. Extravasation Identification. Whole body Planar images (left) and RAVIN Cam (Right).

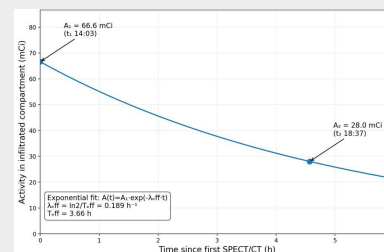


Figure 3. Extravasated Activity clearance from quantitative SPECT/CT (two time-points).

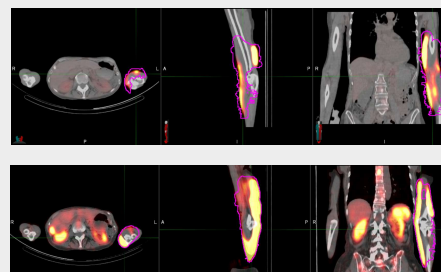


Figure 4. Segmentation and quantification of the extravasation using xSPECT/CT. The upper panel shows the subcutaneous distribution of the extravasated activity in the arm at the first time point, while the lower panel shows the whole-body redistribution of activity at the second time point.

RESULTS

- Quantitative SPECT/CT estimated 66.6 mCi of infiltrated activity, in close agreement ($\approx 7\%$) with the handheld M3D CZT detector (61.8 mCi)
- Serial SPECT/CT imaging over 4.57 h demonstrated rapid clearance, with activity decreasing from 66.6 mCi to 28.0 mCi and yielding an effective half-life (T_{eff}) of 3.66 h, consistent with literature values for rapid local clearance ($\sim 3-4.25$ h) [3,4]
- Patient-specific dosimetry (MIRDcalc sphere model, self-dose) calculated a mean absorbed dose of 1.04 Gy to the 1070 mL extravasated soft-tissue compartment (represents average dose to the defined infiltrated volume).
- Two low-dose CT scans acquired for localization and attenuation correction contributed a total effective dose of 15 mSv ($CTDI_{vol} = 5.37$ mGy each), as estimated with MIRDcalc [2].
- The short measured T_{eff} did not increase patient-release restrictions; the site default assumption of 56.1 h remained more conservative than the case-specific clearance kinetics.

CONCLUSIONS

- This case Demonstrates a practical, clinically feasible workflow for Lu-177 extravasation management using portable detectors and quantitative SPECT/CT.
- Enables rapid identification of extravasation with a handheld CZT detector, followed by accurate activity quantification (~ 66 mCi in 1070 mL soft-tissue volume).
- Serial SPECT/CT at a second time point allows effective half-life estimation ($T_{eff} = 3.66$ h) and patient-specific self-absorbed dose calculation
- Despite infiltration of $\sim 1/3$ of the planned activity, rapid clearance limited mean compartment dose to ~ 1 Gy
- Workflow adds only low CT dose burden while supporting standard patient-release criteria.
- Future work will refine tissue-at-risk dosimetry (skin/subcutaneous layers) and correlate results with clinical outcomes

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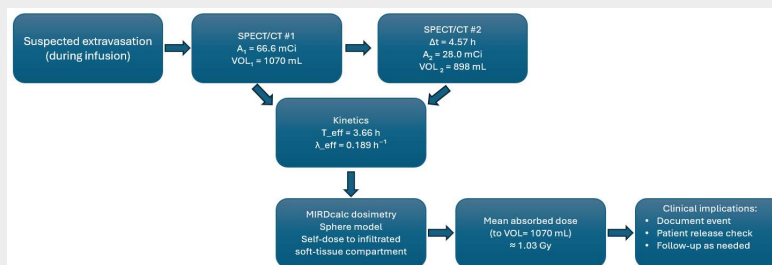


Figure 1. SPECT/CT-based extravasation dosimetry workflow