



SERVIZIO SANITARIO REGIONALE  
EMILIA-ROMAGNA  
Azienda Unità Sanitaria Locale di Reggio Emilia  
IRCCS Istituto in tecnologie avanzate e modelli assistenziali in oncologia



# Comparison between a convolution based and Monte Carlo based dosimetry software

D. Finocchiaro , S. Berenato, E. Grassi, F. Fioroni, G. Castellani, N. Lanconelli, A. Versari, E. Spezi, M. Iori



# Aim

- In this work we compared two systems for dosimetry in MRT that use different techniques of calculation:



$$\bar{D}_{(voxel_k)} = \sum_{h=0}^N \tilde{A}_{(voxel_h)} \times S_{(voxel_k \leftarrow voxel_h)}$$

$$S_{(voxel_k \leftarrow voxel_h)} = \sum_i \Delta_i \cdot \frac{\phi_i(voxel_k \leftarrow voxel_h)}{m_{voxel_k}}$$

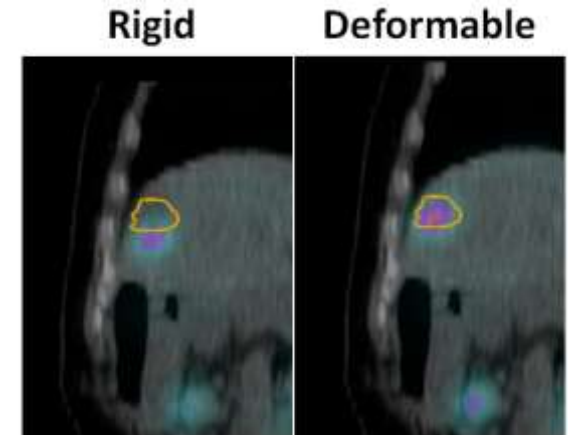
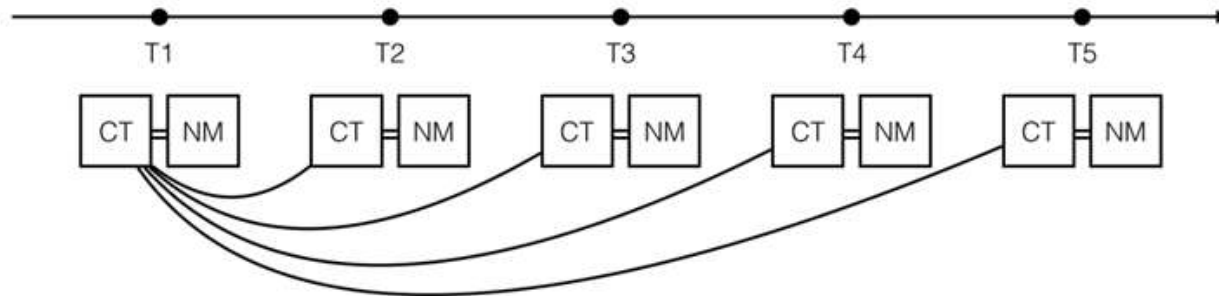
$$D = \left[ \sum_{i=1}^{N-1} (\dot{D} \cdot \tilde{A}_{total}^{i \rightarrow i+1}) \right] + [\dot{D}^N \cdot \tilde{A}_{total}^{tail}]$$

- The aim is to assess how calculation modality impacts on dosimetry results.

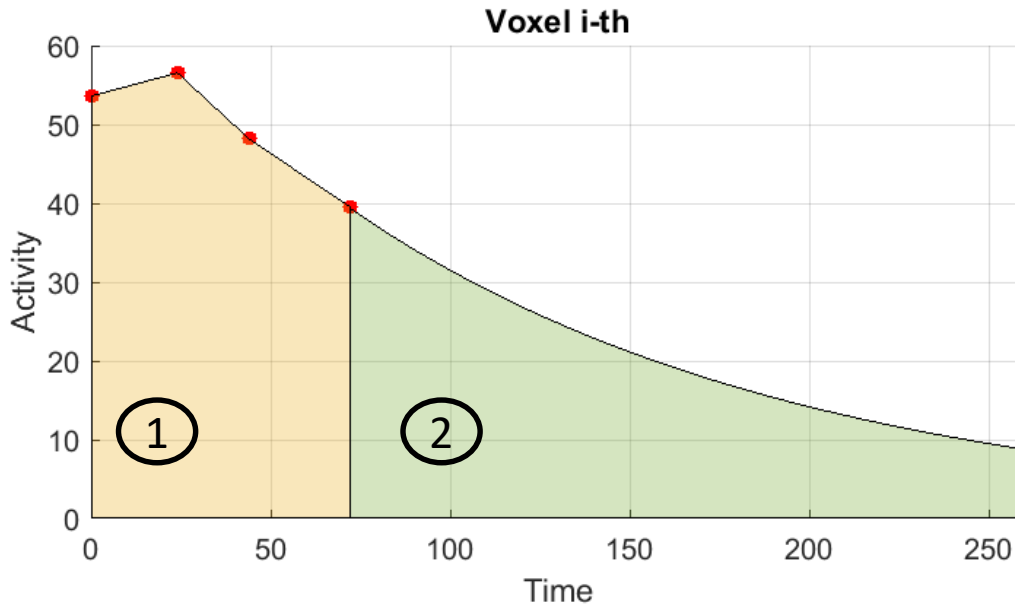
# Material and methods

Study on **20 patients** enrolled in a PRRT clinical trial:

- Dosimetry at the first course of therapy after a therapeutic administration of  **$^{177}\text{Lu}$ -DOTATOC**.
- **5 SPECT/CT** scans with a Siemens Symbia T2 gamma camera at 1, 4, 24, 44, 72 h p.i.
- SPECT images were aligned to the first CT image using a **deformable registration** with the Velocity console (Varian Medical System, USA).



Both VoxelMed and RAYDOSE perform integration of Time-Activity curves using Trapezoidal and Analytical methods:



① Integration by **Trapezoidal method of voxel activity**

② Extrapolation by **whole organ activities**

VoxelMed



Bi-exponential:

$$Ae^{-a \cdot t} + Be^{-b \cdot t}$$

RAYDOSE



Mono-exponential:

$$Ae^{-a \cdot t}$$

- The **same set of images** and the **same VOI** were the starting point for dose calculation with both the software
- Absorbed doses were calculated for

**Organs:**

Left kidney  
Right kidney  
Liver  
Spleen

**Tumours:**

Max 4 tumours per patient  
Total 23 tumours were considered



21 out of 23 were hepatic lesions  
(1 pancreatic and 1 lymph node)

- For the comparison RAYDOSE was considered as the reference and **Error (%)** of VoxelMed mean absorbed dose was calculated as follows:

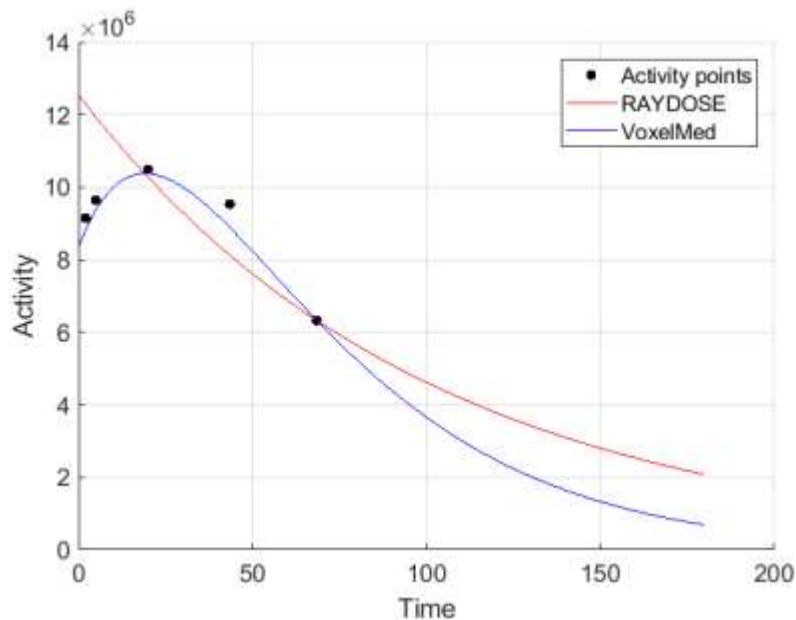
$$Error (\%) = \frac{VoxelMed - RAYDOSE}{RAYDOSE} \cdot 100$$

Furthermore **DVH** and **dose maps** were qualitatively compared, and the **Lin's concordance coefficient** was estimated

To strictly evaluate differences in calculation modality, comparison was also performed using the same **TIAC**:

Calculation with **VoxelMed** was repeated using **the same k** (the effective decay constant) used for RAYDOSE calculation.

Different  
fitting curves:



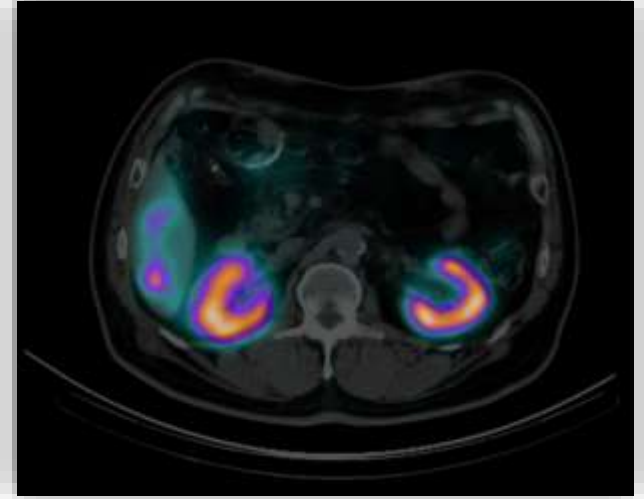
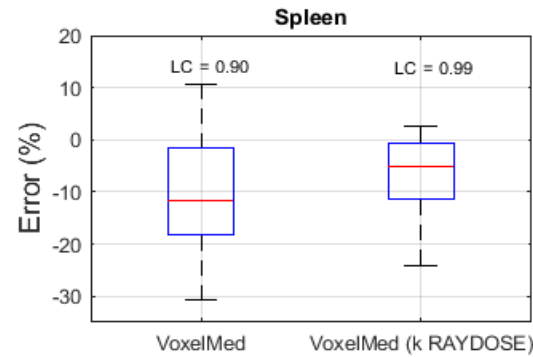
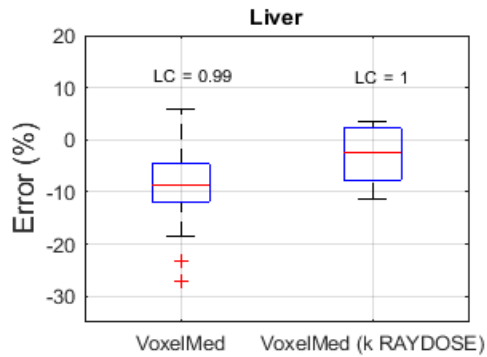
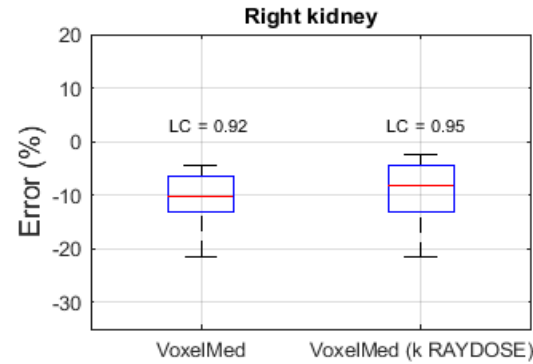
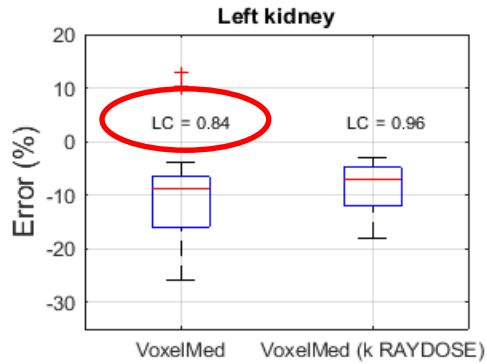
Comparisons were performed between:

**VoxelMed vs RAYDOSE**

&

**VoxelMed (k RAYDOSE) vs RAYDOSE**

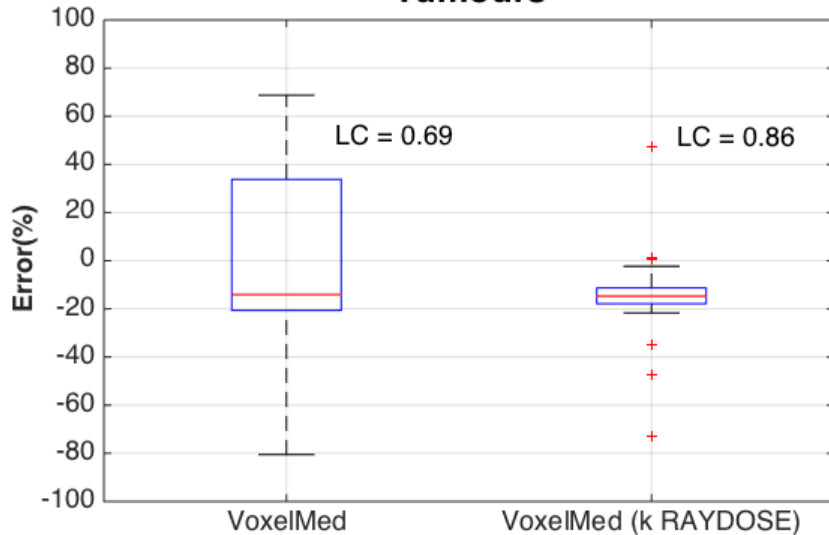
# Results



## Mean Error (%)

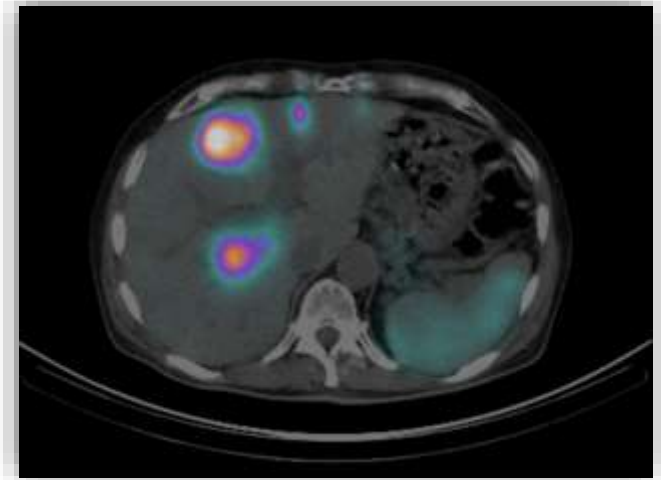
Organ	VoxelMed	VoxelMed (k RAYDOSE)
Left kidney	-10,8	-8,3
Right kidney	-8,6	-9,2
Liver	-7,9	-0,9
Spleen	-6,1	-5,8

## Tumours



- For 3 tumours out of 23 Error (%) > 100%
- These differences were reduced when the same k was used
- Less variability and higher correlation was observed when the same k was used
- In case of VoxelMed (k RAYDOSE) Error(%) < 0 for all the lesions (excluded outliers)

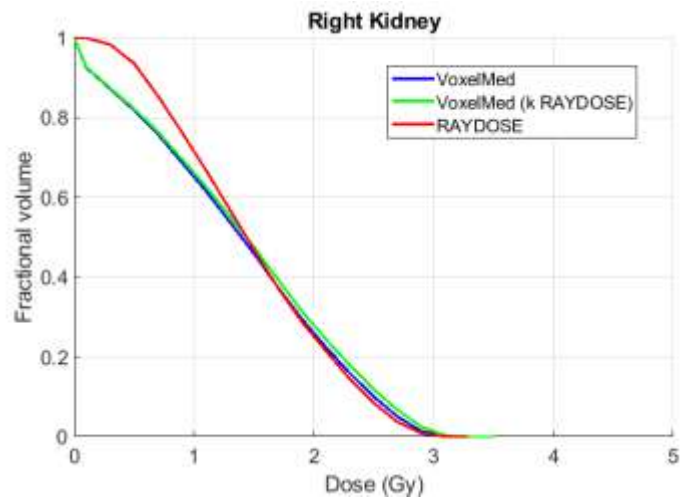
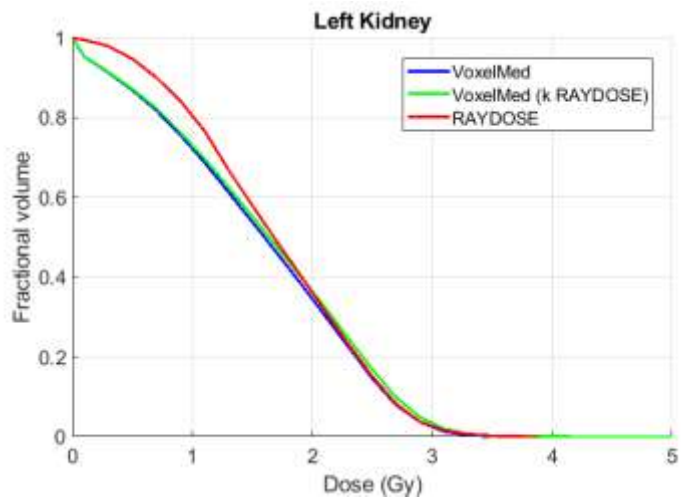
	VoxelMed	VoxelMed (k RAYDOSE)
Mean	-11,5	-16
Median	-17,6	-14,8
25 <sup>th</sup> perc	-22,0	-19,0
75 <sup>th</sup> perc	30,8	-11,8





Finally we qualitatively compared the DVH and Dose maps:

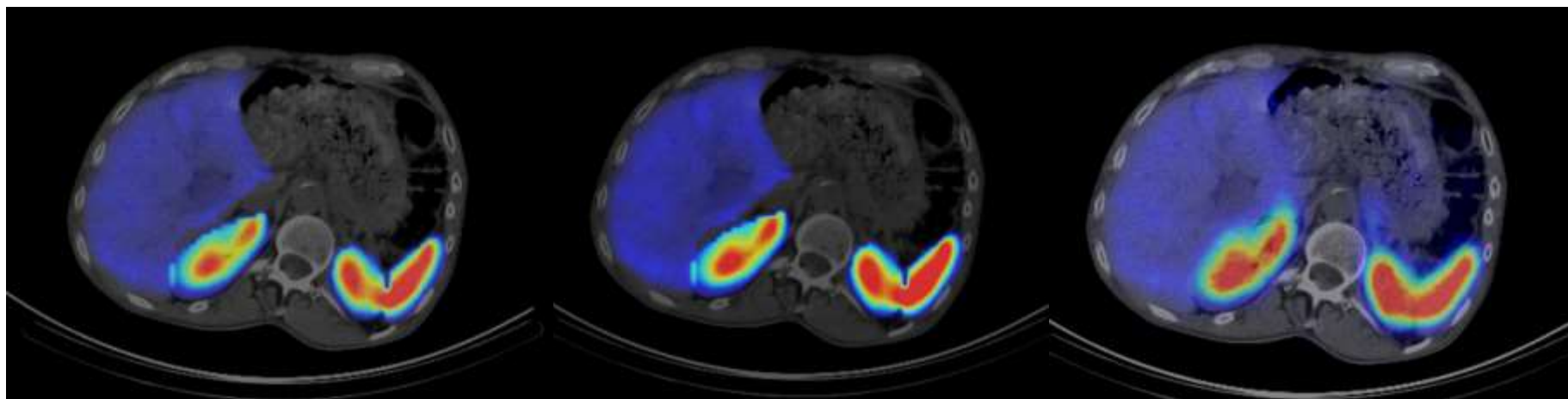
Example of a patient with typical Error (%)



VoxelMed

VoxelMed (k RAYDOSE)

RAYDOSE



# Conclusion

- A general underestimation of absorbed doses calculated with VoxelMed was observed, as compared to RAYDOSE: average -9% error was observed for organs, while -18% for lesions.
- Concordance between VoxelMed and RAYDOSE was good for organs ( $LC > 0.84$ ), while greater differences were observed for tumours ( $LC = 0.62$ ).
- Differences were reduced when the same  $k$  was used: -6% for organs and -15 for lesions. Also concordance was largely increased ( $LC > 0.95$  for organs and 0,86 for lesions).
- In conclusion convolution methods allows to perform high accuracy dosimetric calculations with reduced computational time.
- Particular attention must be paid to the activity fitting that could have a large impact on results.

**Thank you for  
your attention**