

# Measurements of dose point kernels using GATE Monte Carlo toolkit for personalized convolution dosimetry

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# Outline

- Introduction
- Objectives
- Methodologies
- Results
- Conclusion

# Introduction

- ✓ Dose Point Kernels (DPKs):
  - are defined as energy deposition profiles from a point isotropic sources
  - method to compute the absorbed dose from the non-uniform activity or high gradient activity distributions
- ✓ Usefulness of kernels?
  - Convolution dosimetry: calculate the dose distributions using the convolution of 3D-dose kernel matrix with cumulated activity map furnished by quantitative SPECT/CT or PET/CT images.
  - Cumulated activity is represented by area under the time activity curve
- ✓ Ultimate goal: patient-specific dosimetry

# Patient-specific dosimetry:

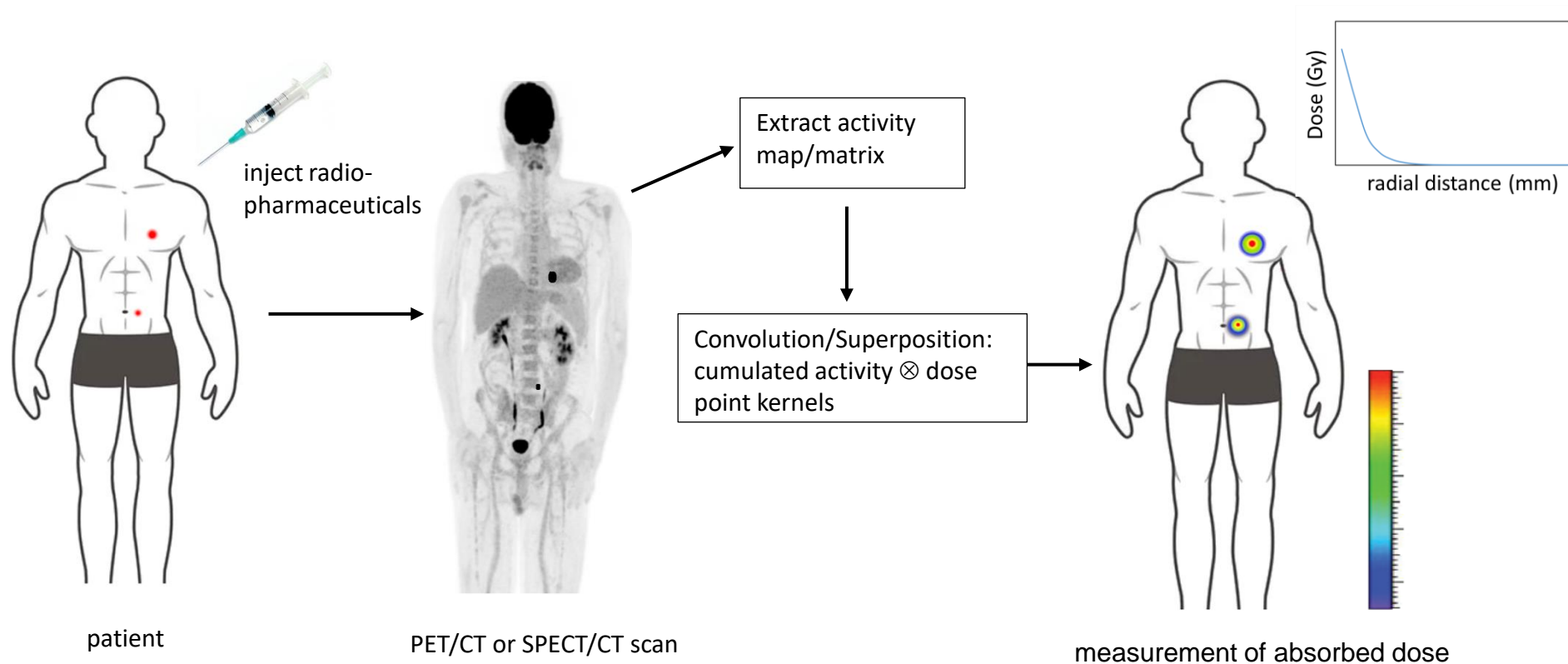


Figure: Schematic representation of the workflow for the patient-specific dosimetry using the Monte Carlo generated dose point kernels and the patient PET/CT scan.

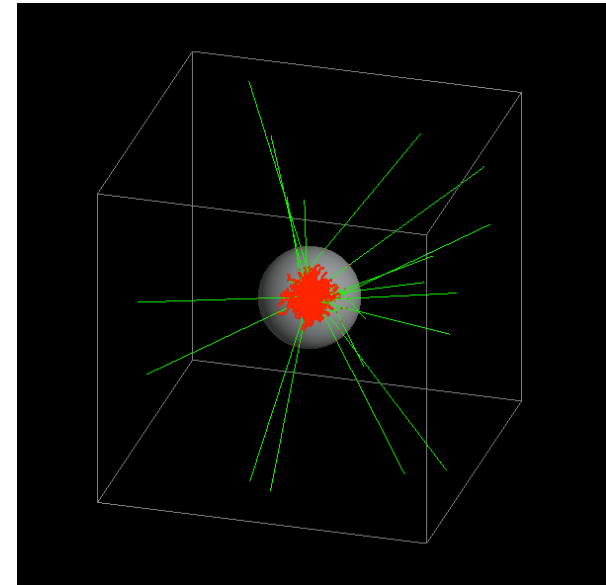
➤ Because of its complex nature it has not been implemented in clinical settings yet!

# Monte Carlo approaches to generate dose point kernels

- ✓ approx. solution of transport equations, no exact/non-trivial analytic solutions
- ✓ simulation of stochastic process ( $e^-$  s and  $\gamma$ -s transport)
- ✓ Geant4, GATE, EGS, FLUKA, PENELOPE, ETRAN, MCNP, SMOOPY etc.
- ✓ GATE: Geant4 Application for Tomographic Emission
- ✓ GATEv8.1 and v7.2 were used
- ✓ *electronStepLimiter*, production threshold cuts

# Simulation mediums

- Dose point kernels algorithms are based on an idea of homogeneous medium
- Human body is composed of different tissues: water, bone, blood, lung, adipose, red marrow ...
  - $\rho$  (g/cm<sup>3</sup>)
  - elemental composition ( $Z_{\text{eff}}$ )



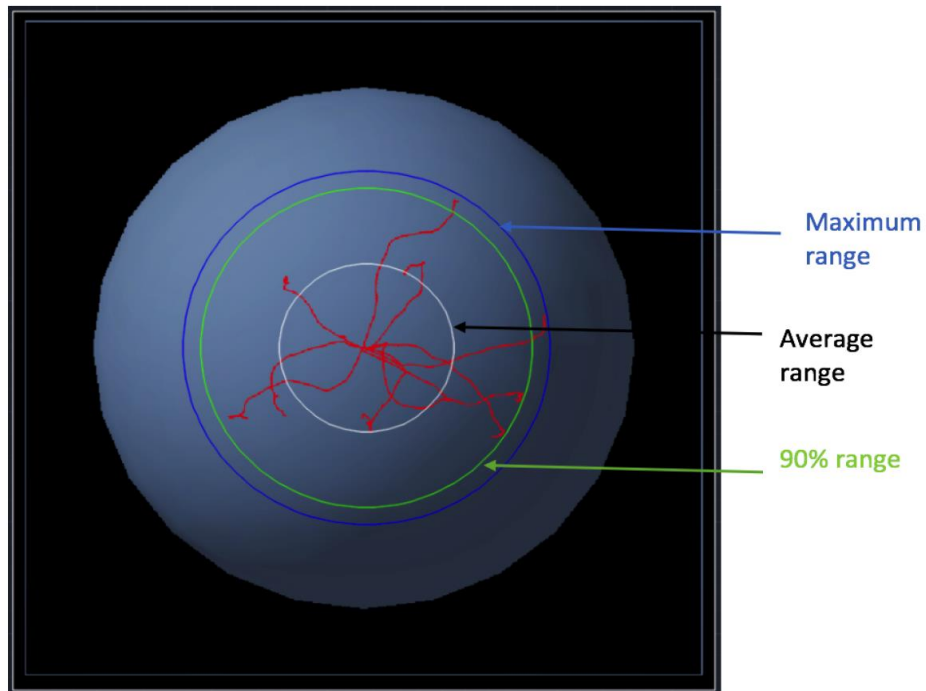
Materials	bone	blood	lung	water	red marrow	adipose
$Z_{\text{eff}}$	11.87	7.78	7.74	7.42	7.21	6.47
$\rho$ (g/cm <sup>3</sup> )	1.85	1.06	0.26	1.00	1.03	0.92

We want to have the DPKs for various tissues to find whether it depends on tissue type.

# Methodologies

- Monoenergetic electron DPKs:

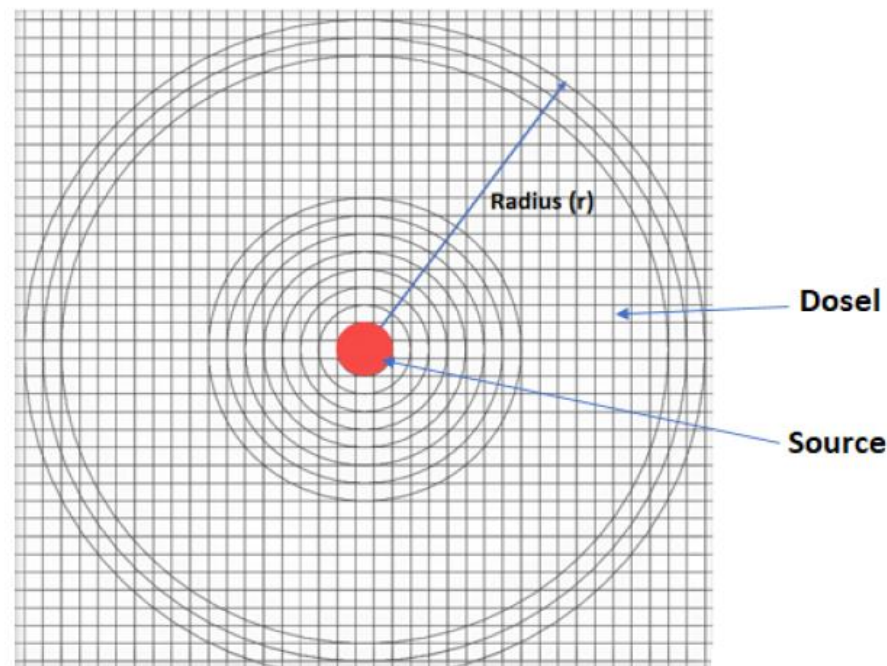
$$J(r/R_{\text{CSDA}}) = 4\pi r^2 D(r, E) \frac{R_{\text{CSDA}}}{E_0}$$



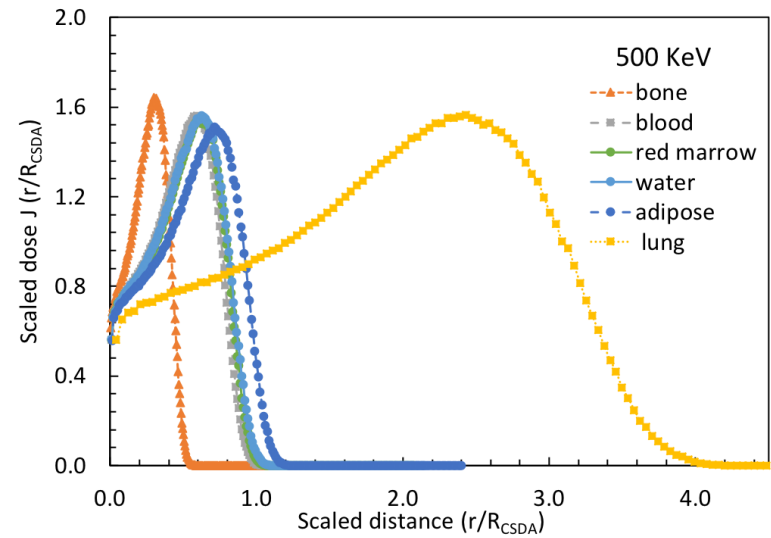
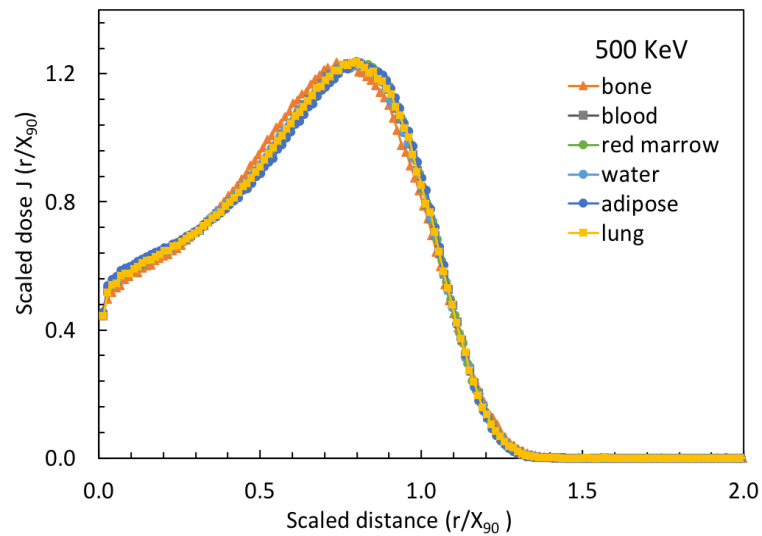
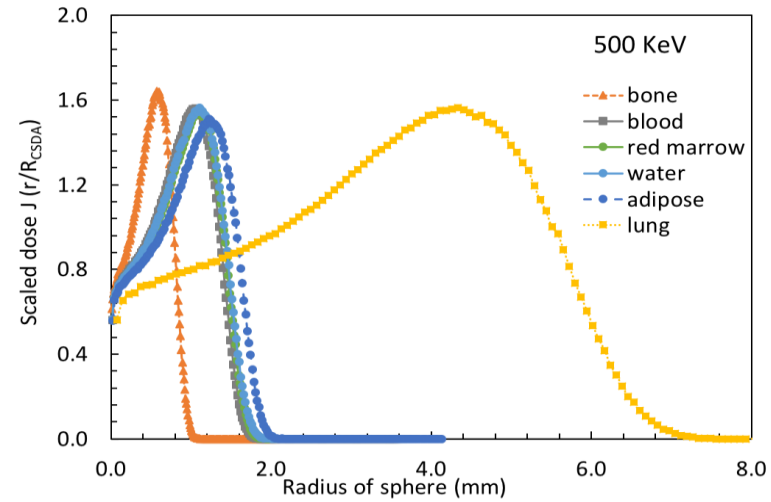
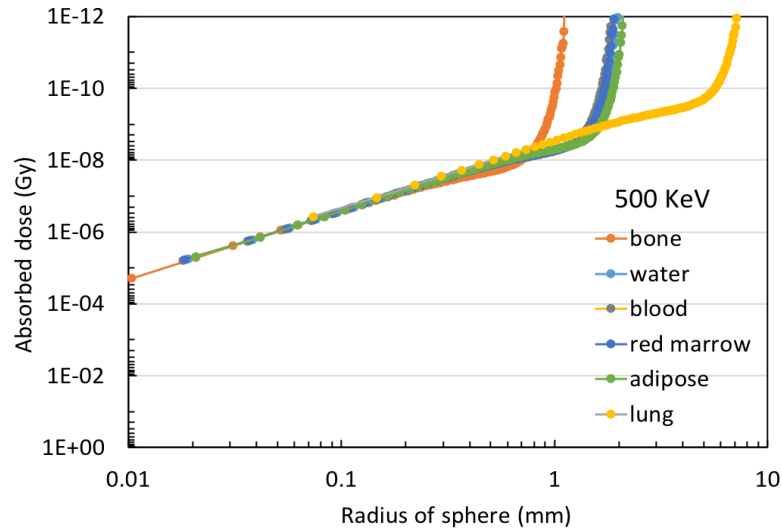
- Beta spectrum DPKs:

$$J(r/X_{90}) = 4\pi r^2 D(r, E) \frac{X_{90}}{E}$$

$$X_{90} = 4\pi\rho \int_0^{X_{90}} r^2 \Phi(r, E) dr = 0.90$$

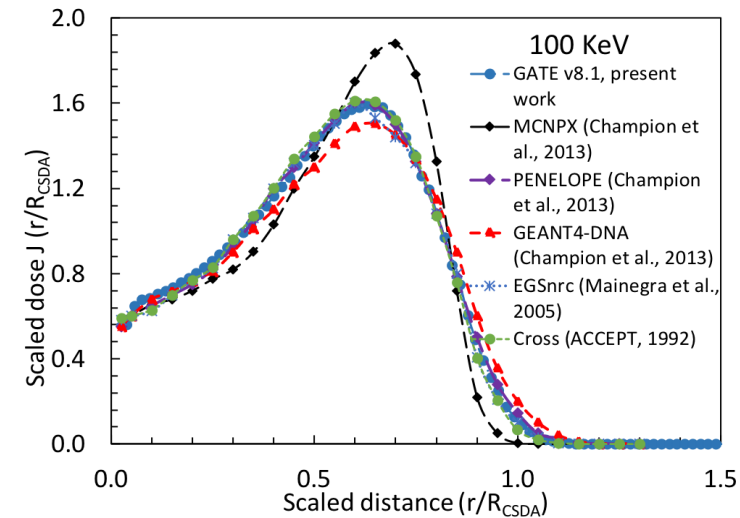
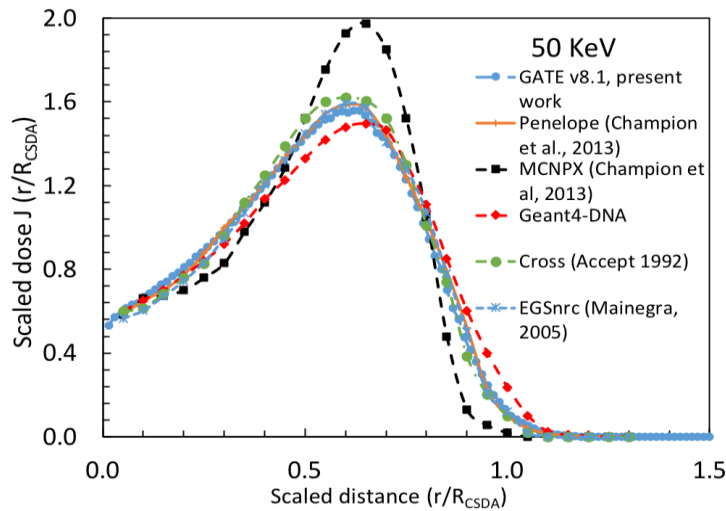
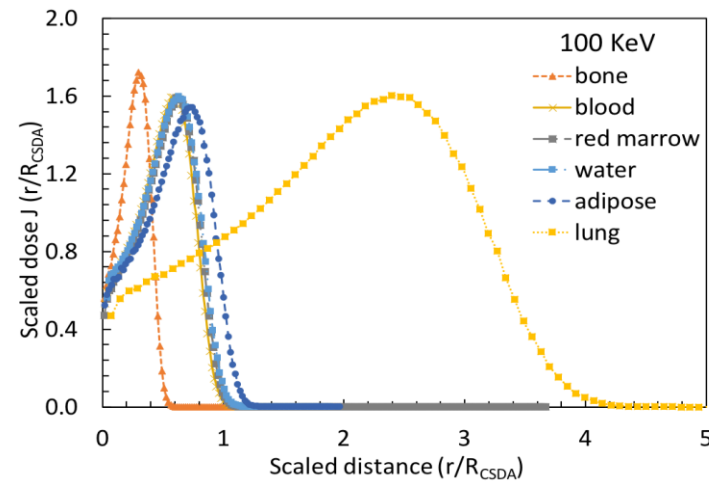
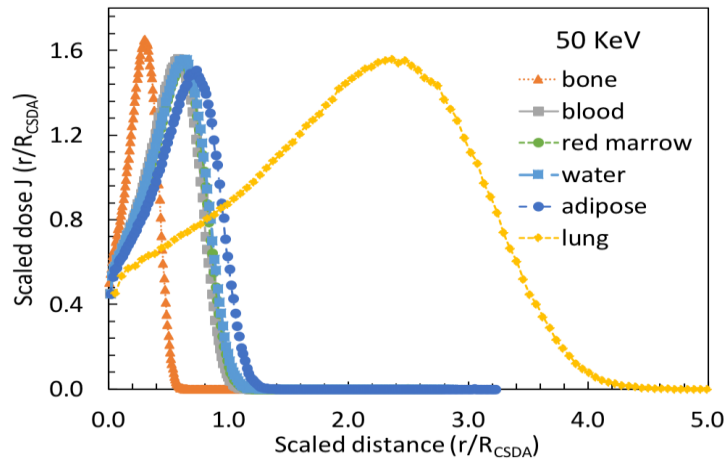


# Results: (i) Monoenergetic electrons dose point kernels: 500 KeV

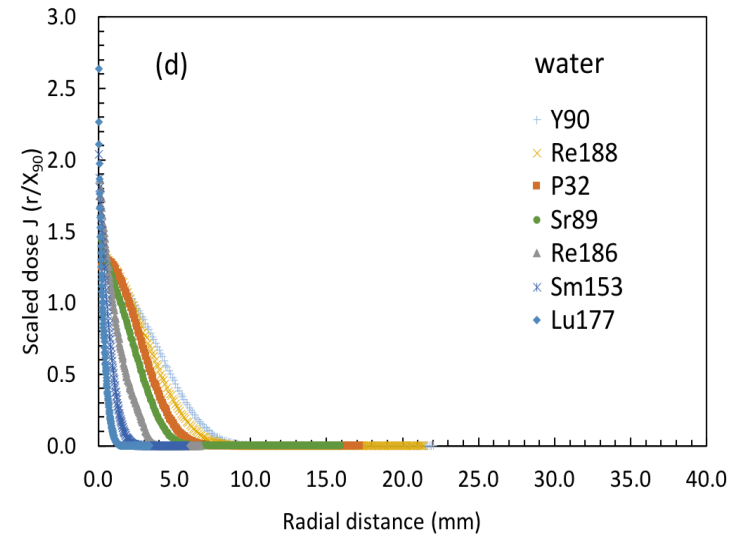
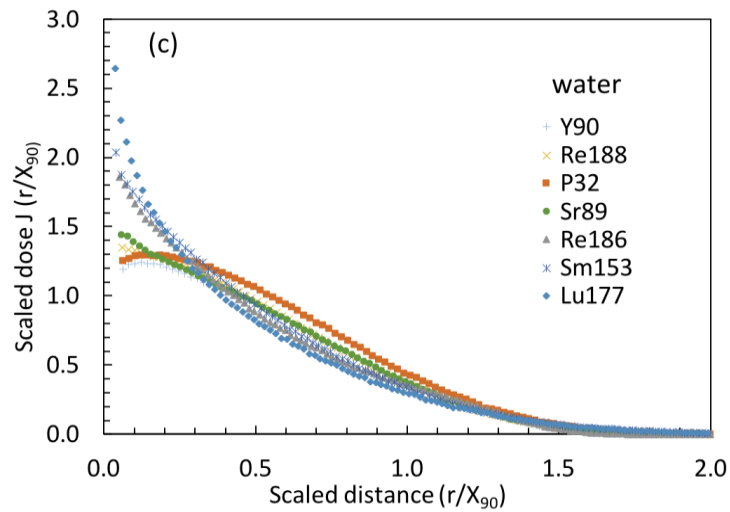
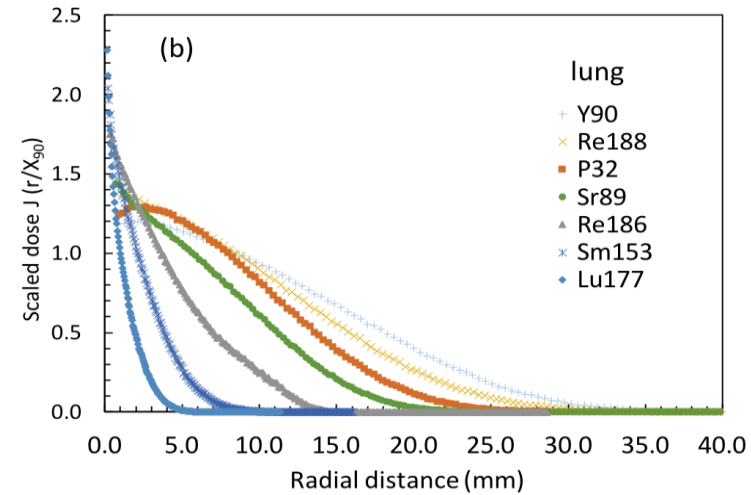
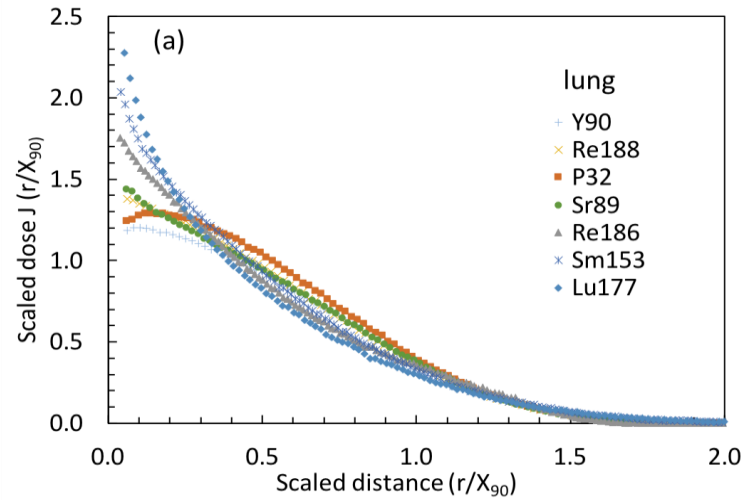




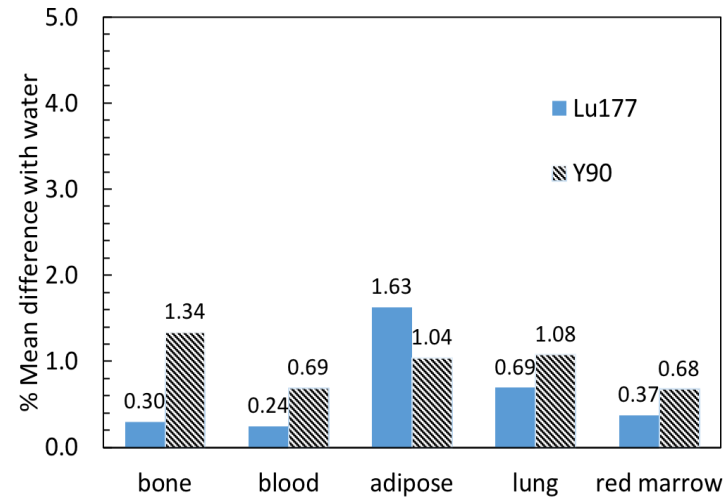
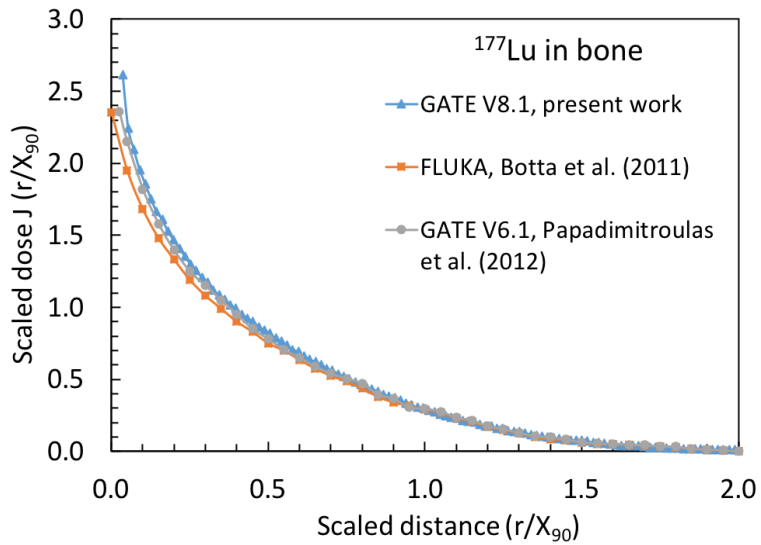
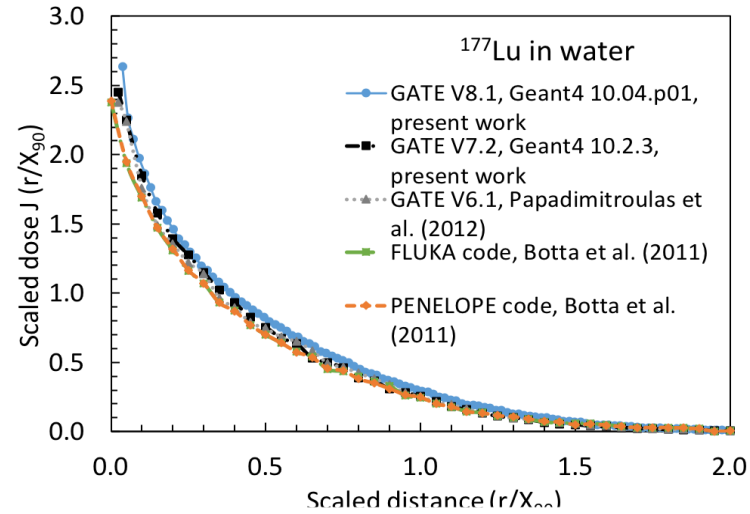
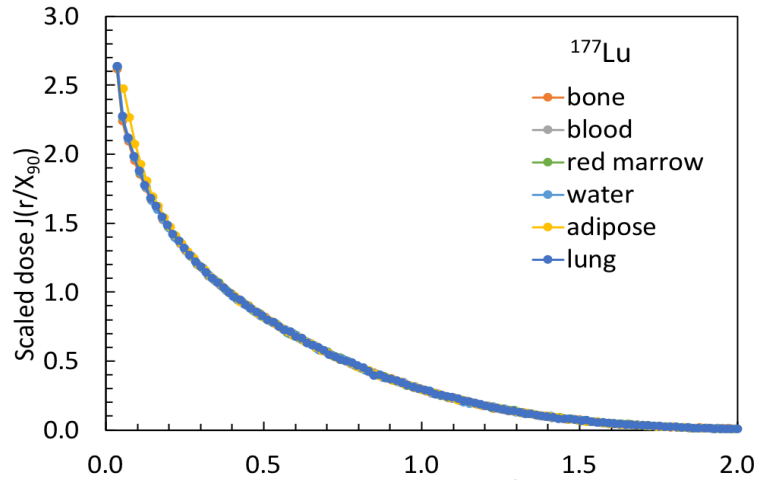
# Comparison of monoenergetic electrons dose point kernels:



# Results: (ii) beta radionuclides dose point kernels



# Impact of tissue type on dose point kernels?



# Conclusion

- ✓ Dose point kernels of monoenergetic electrons (0.015-10) MeV and radionuclides beta emitters has been simulated
- ✓ Minimal discrepancies are observed between water and other tissues kernels when scaled with  $X_{90}$  for all simulated isotopes
- ✓ Impact of tissue type has been found to be minimal for purposes of dosimetry
- ✓ Use of single kernel generated in water may be sufficient for 3D dose calculations
- ✓ Future work is targeting experimental validation of dose point kernels

# Experimental validation

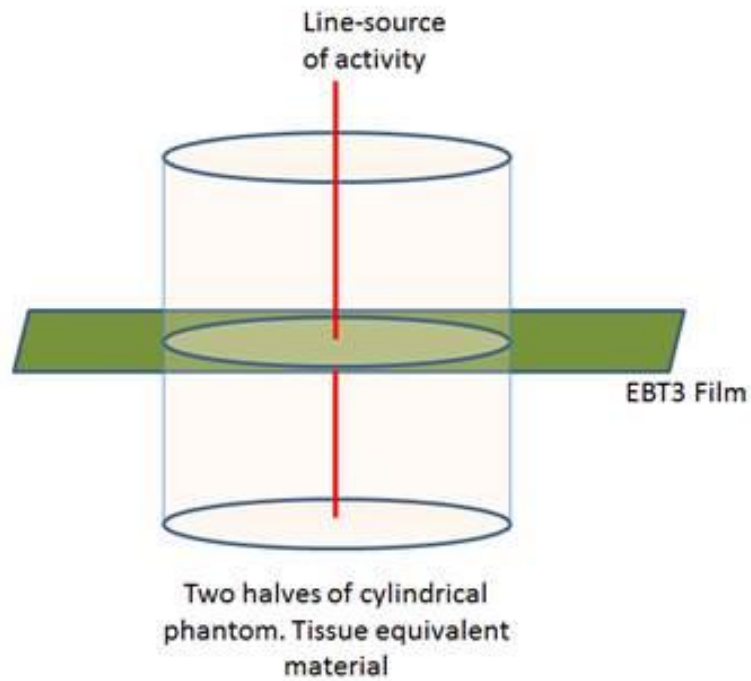


Fig 1: detector geometry with Gafchromic films

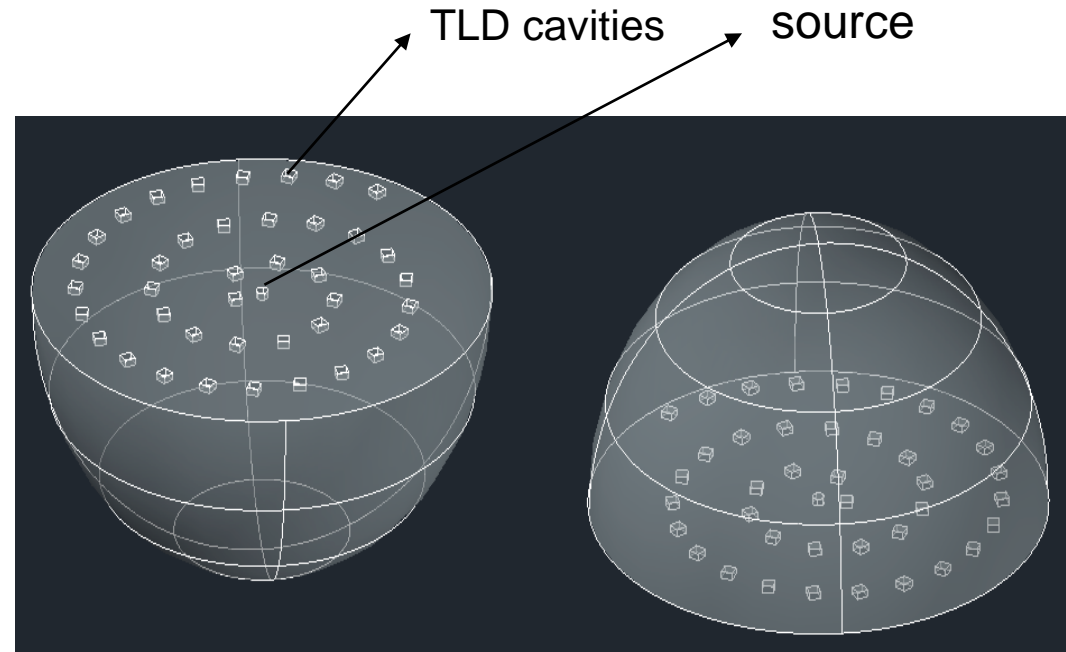


Fig 2: detector geometry equipped with TLDs