

The Italian project for a proton Computed Tomography (pCT) device

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and the PRIMA collaboration

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1. Why the pCT?

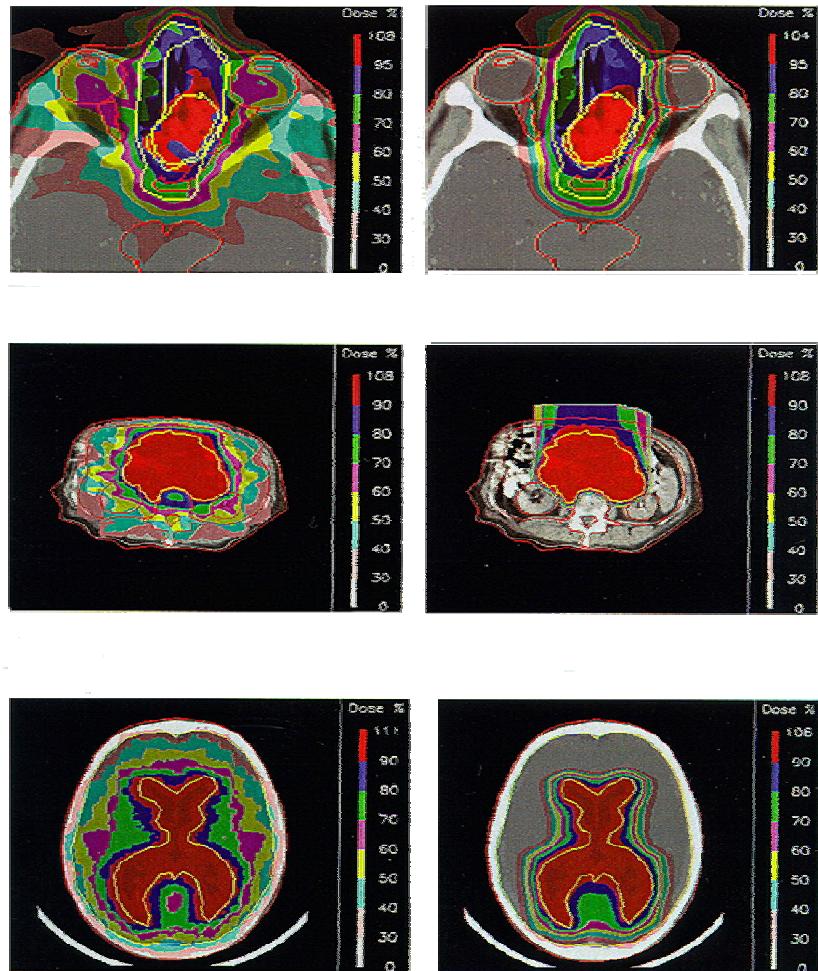
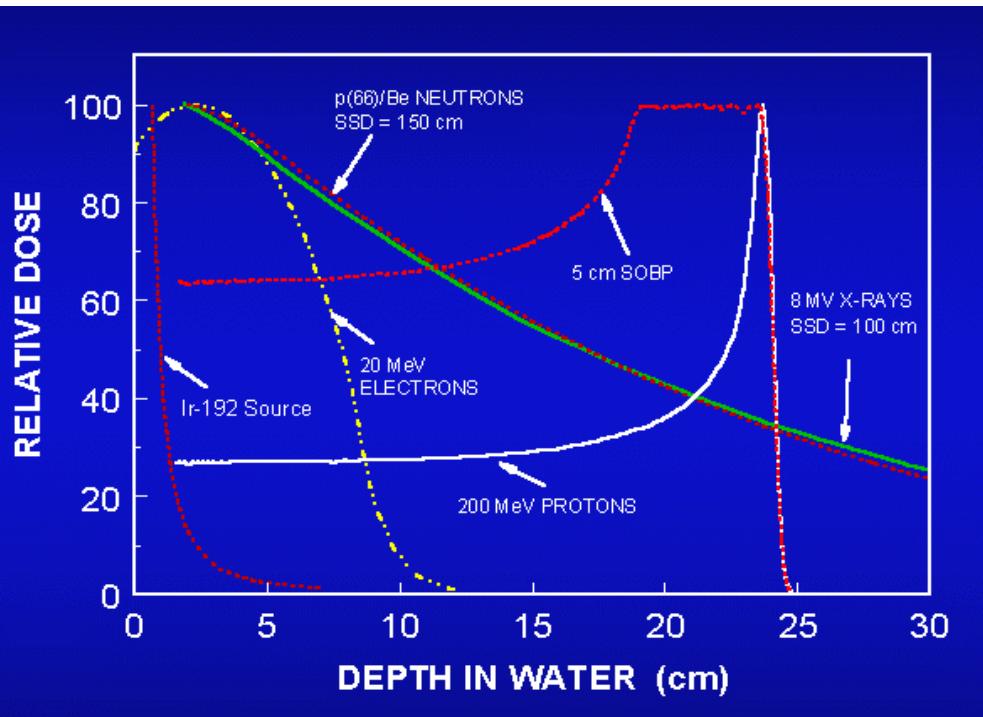
2. The proton Computed Tomography: history, physical principles and status of the art

3. Design of the detector

4. Role of Monte Carlo simulation

WHY A pCT DEVICE ?

Proton therapy is a growing radiation treatment technique (more 35 centers today)



MAIN ISSUES IN PROTON THERAPY QUALITY

- ❑ Patient positioning

- ❑ Dose planning

Actually TPS are based on the xCT images as input and this bring a sensible amount of imprecision

DOSE PLANNING IN PROTON THERAPY

The *stopping powers* represent the main parameter for the dose calculation in a proton treatment planning

In proton therapy they are indirectly derived measuring and converting, following some calibration curves, the attenuation coefficients μ derived from a conventional CT

THE ERROR INTRINSIC IN THIS CONVERSION

OWN TO THE NOT SIMPLE DEPENDENCE OF μ FROM η_e AND Z OF THE MATERIAL

**IS THE PRINCIPAL FACTOR LEADING TO THE RANGE
INDETERMINATION OF PROTON**

Studies by PSI researchers on phantom and animal models demonstrated that the used conversion

$$\mu \leftrightarrow \text{stopping power}$$

and with actual calibration model produce an average indetermination on proton range of 3%

(up to 10 mm in the head)

Range uncertainty



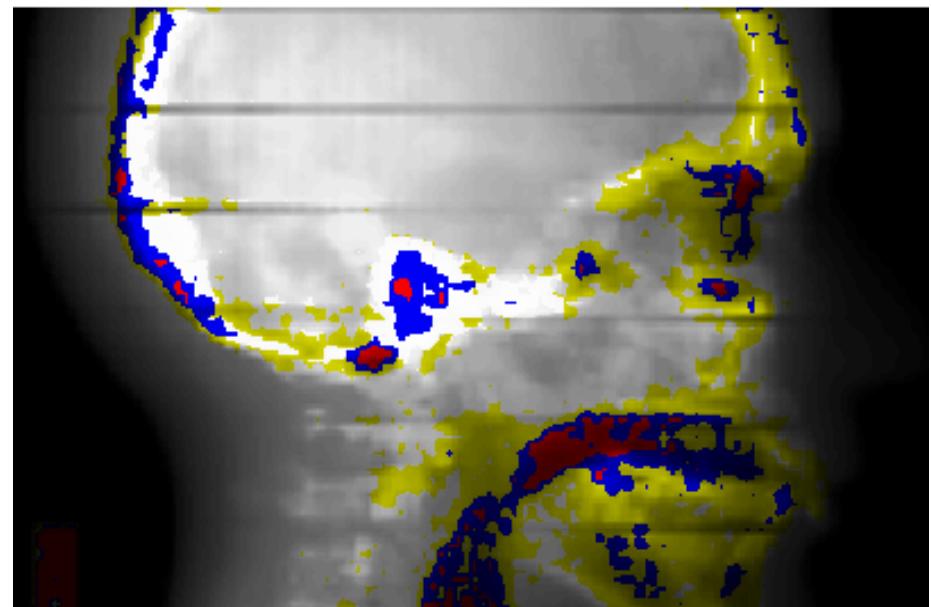
> 5 mm



> 10 mm



> 15 mm



Alderson Head Phantom

Schneider U. (1994), "Proton radiography as a tool for quality control in proton therapy," Med Phys. 22, 353.

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Already in his Nobel lecture, for the computed tomography realisation, Allan Cormack shown his studies carried out with Andreas Koehler and published on J. of Appl. Phys. nel 1963 e 1976, on the possibility of a protonComputed Tomography



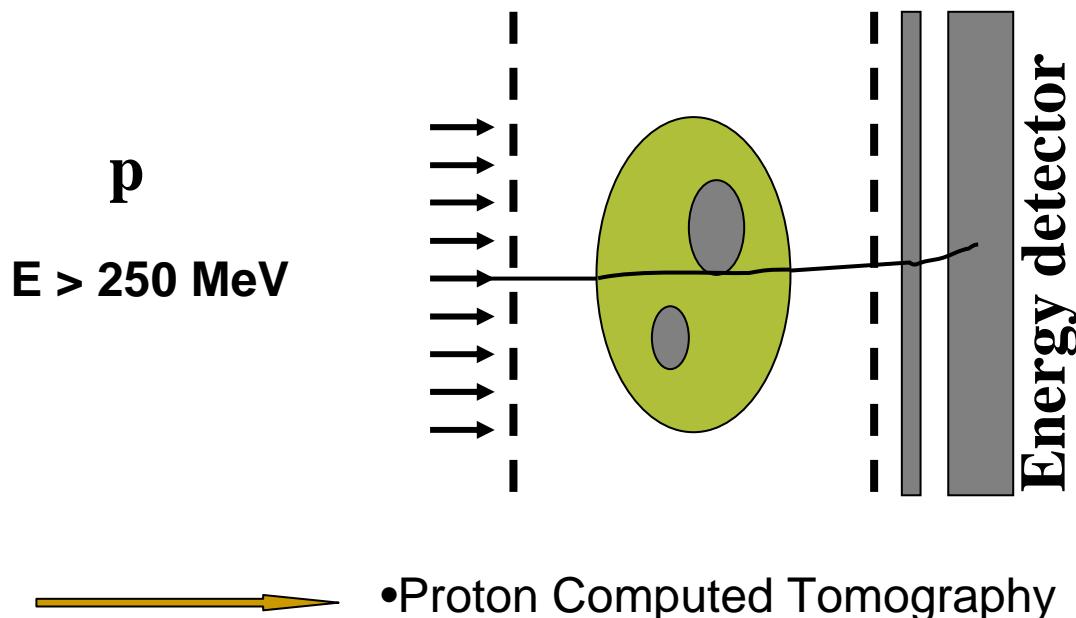
<< Why use proton? First there is the question of dose..... There is an agreement with theoretical calculation that the dose required for proton scans is five to ten times less than for X-Ray scans for the same amount of informations.>>

<< Second: due to the different mechanism of interaction with matter one ought to see different things using different radiations.....>>

TWO MAIN REASONS STOPPED THE INTEREST OF PCT IN THE NEXT YEARS:

- A VERY SMALL NUMBER OF PROTON-THERAPY CENTERS
- THE IMPOSSIBILITY TO REACH SUFFICIENT LEVELS OF SPATIAL
AND DENSITY RESOLUTION OWN THE INTRINSIC PRESENCE OF
THE COULOMBIAN MULTIPLE SCATTERING

METHOD EQUIVALENT TO THE X-RAYS IMAGING BUT HERE THE MAIN INFORMATION IS ENERGIES OF TRAVERSING PROTONS DESPITE ATTENUATION COEFFICIENTS OF PHOTONS



Knowledge of entry and exit energy of a proton allow the estimation of the integrated (projected) relative electron density along a path L

The mean rate of energy loss is given by the Bethe-Bloch equation, written in a useful for pCT:

$$-\frac{dE}{dx}(r) = \eta_e(r)S[I(r), E(r)]$$

$$\int_L \eta_e(\vec{r}) d\vec{r} = K \int_{E_{out}}^{E_{in}} \frac{dE}{S(E)}$$

Integrated electron density

Inverse of Stopping Power

Like in xCT the pCT image reconstruction is the inversion of right integral for the unknown electron density distribution

BUT

the exact path L in the pCT is unknown due to Multiple Coulomb Scattering and only initial and final information can be measured for each proton

MAIN TASK:

ESTIMATION OF PROTON PATH INSIDE MEDIUM

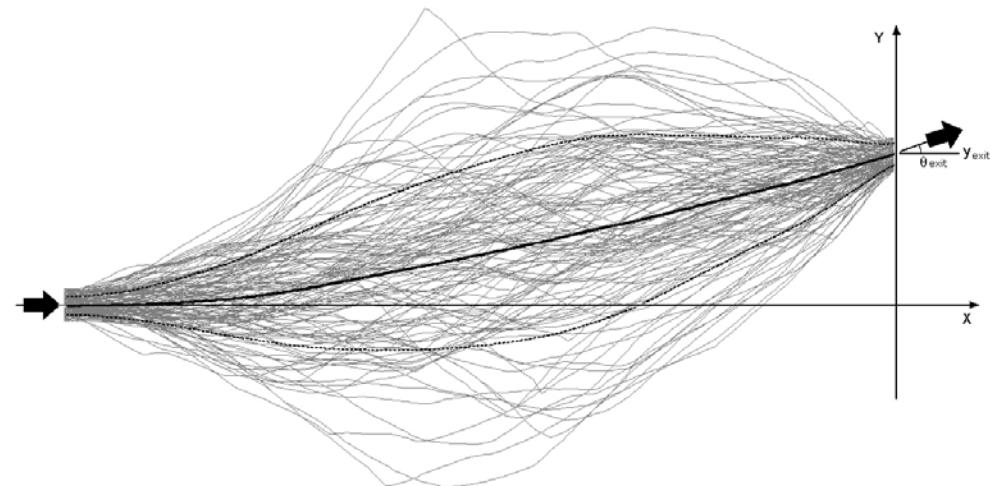
OR BETTER



A MEDIUM PATH

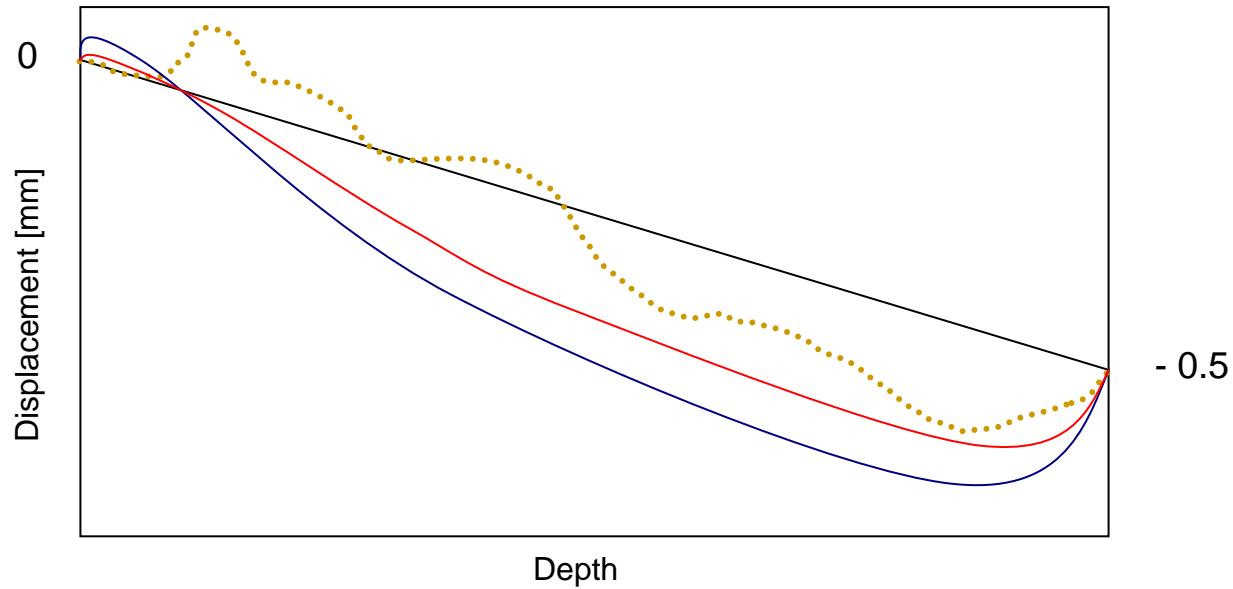
Geant4 simulation of 250
MeV protons in water

THE SINGLE TRACKING
APPROACH



MEDIUM PROTON PATH ESTIMATION

1. Straight-line path
2. Most Likely Path (MLP): from Moliere MCS theory
3. Cubic Spline path
4. Monte Carlo path



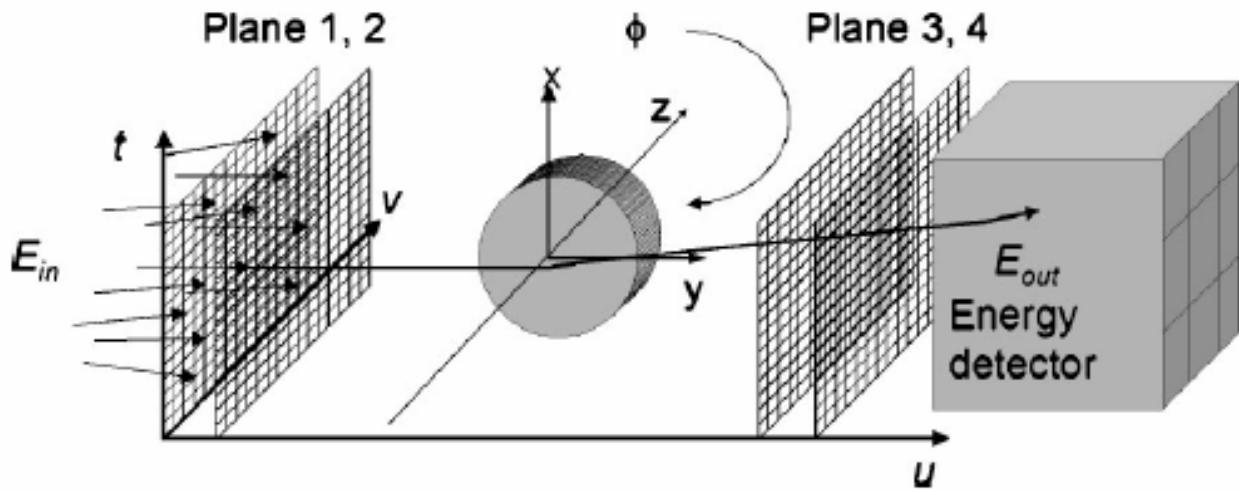
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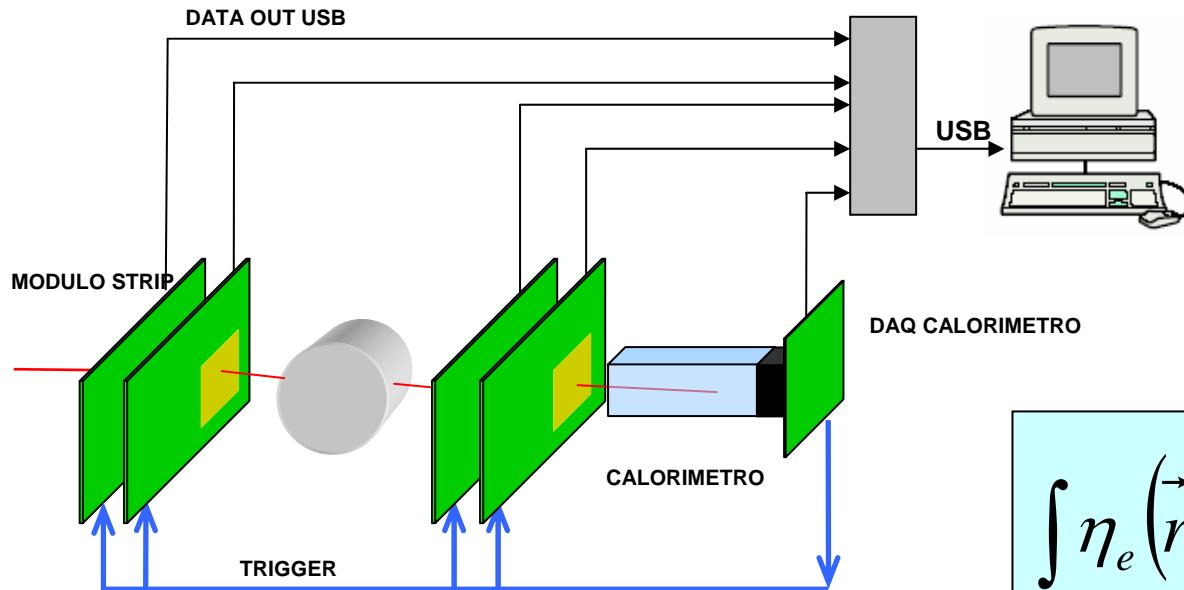
All the studies and prototypes developed in the last years are based on the principle of follow each single proton traversing the medium to investigate



The “single tracking” approach should permit to improve the spatial resolution up to 1 mm or less

$$\int_L \eta_e(\vec{r}) d\vec{r} = K \int_{E_{out}}^{E_{in}} \frac{dE}{S(E)}$$

THE PRIMA DETECTOR



$$\int_L \eta_e(\vec{r}) d\vec{r} = K \int_{E_{out}}^{E_{in}} \frac{dE}{S(E)}$$

Single tracking

Acquisition rate up to 1 MHz

Detector: 2 orthogonal microstrip; 200 um pitch x 256 strips; about 5x5 cm of active area

Trigger from the calorimeter

Requirements for a pCT device

Category	Parameter	Desired Value
Proton source	Energy	≈ 200 MeV (head) ≈ 250 MeV (trunk)
	Energy spread	$\approx 0.1\%$
	Beam intensity	$10^3 - 10^5$ protons/sec
Accuracy	Spatial resolution	< 1 mm
	Electron density resolution	< 1%
Time Efficiency	Installation time	< 10 min
	Data acquisition time	< 5 min
	Reconstruction time	< 15 min (treatment planning) < 5 min (dose verification)
Reliability	Detector radiation hardness	> 1000 Gy
	Measurement stability	< 1%
Safety	Maximum dose per scan	< 5 cGy
	Minimum distance to patient surface	10 cm

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1. Test the estimation of the proton path
2. Evaluate of image quality
3. Evaluate the released dose (as function of image quality!)
4. Evaluate the different reconstruction algorithms

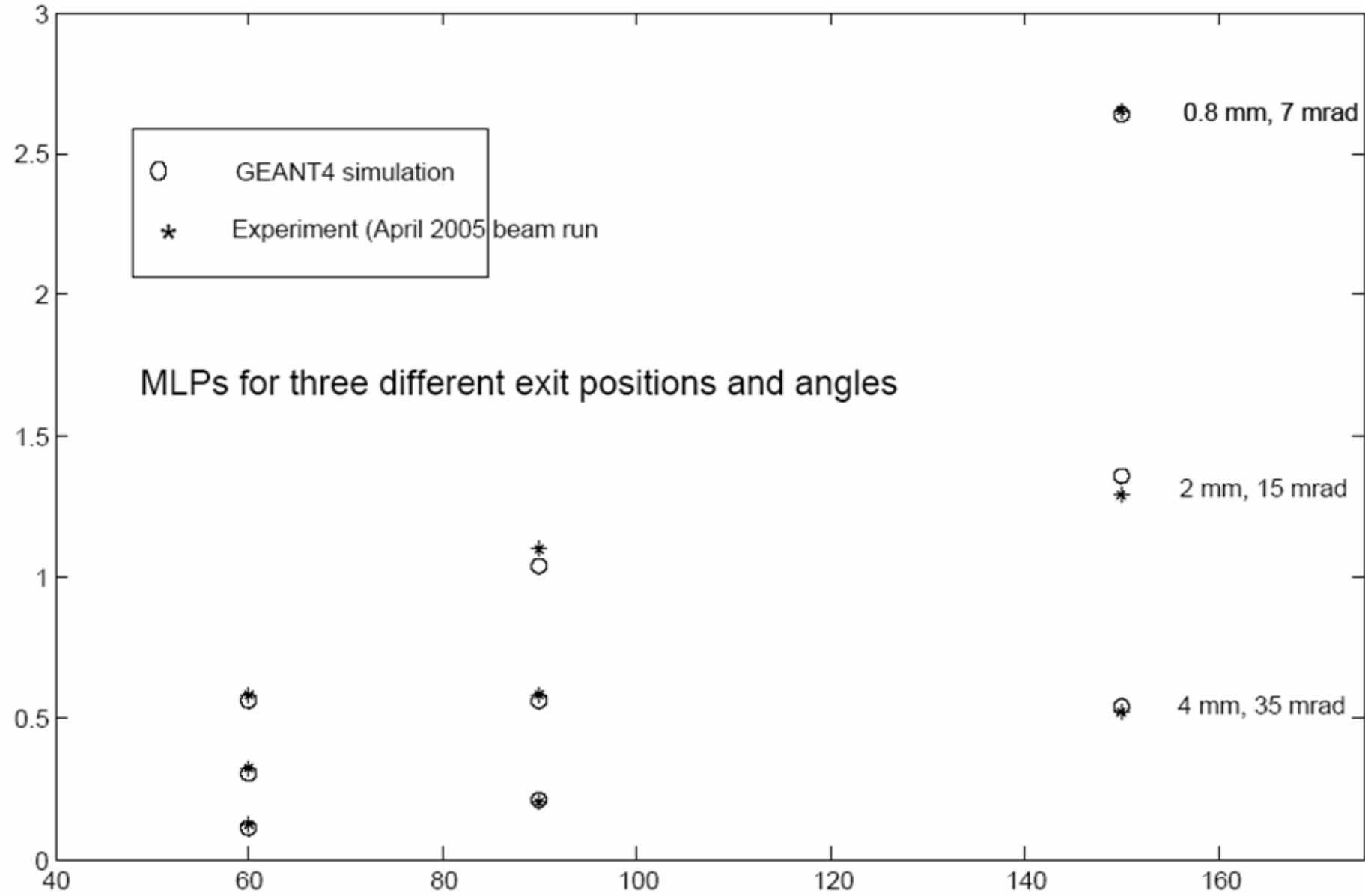
OUR COMPUTATIONAL TOOL

- GEANT4 MC toolkit:
- Electromagnetic processes
- Elastic nuclear scattering
- Inelastic nuclear scattering

First tests of the Williams analytical approach to MLPs

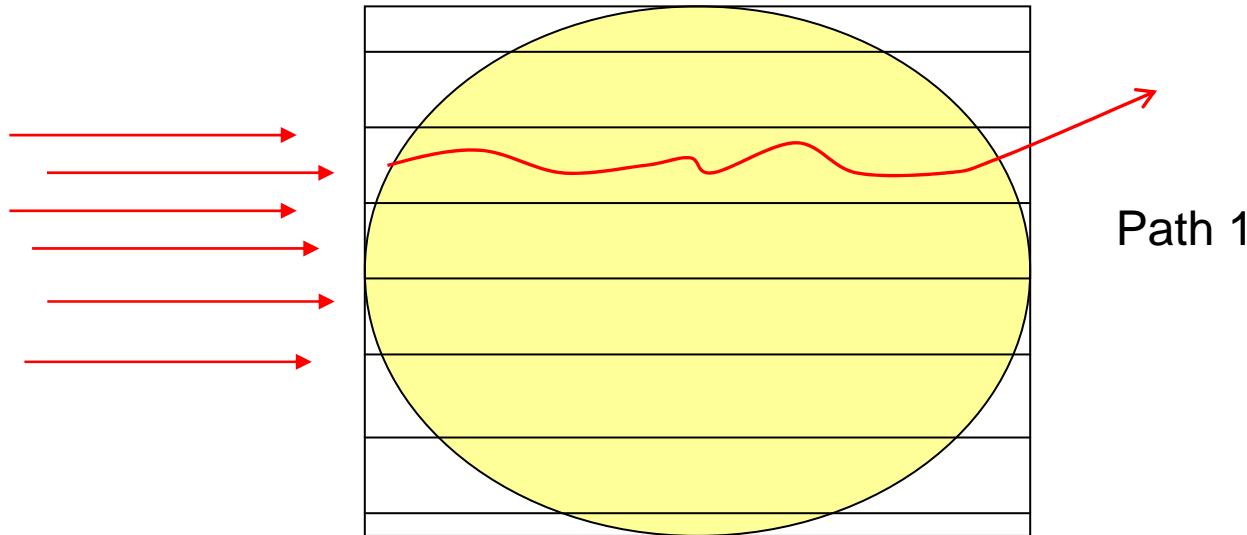


G4 vs Experiment



MLPs for three different exit positions and angles

IMAGE RECONSTRUCTIONS WITH THE FBP ALGORITHM

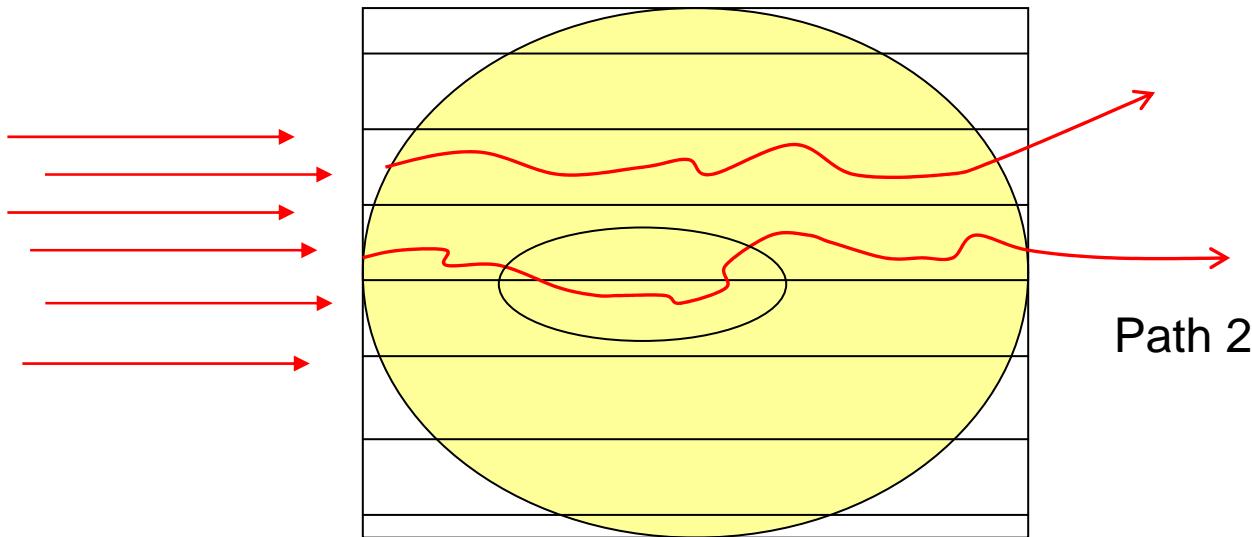


Object is ideally divided in channel and only protons do not exceed the bin AT ANY POINT are considered. Its path can be approximated to a straight line and FBP is applicable

THE ROLE OF MONTE CARLO

A different possibility is consider protons entering and exiting in the same channel but not limit their paths inside it (Path 2). Eventually a constraint on exit angle can be introduced

This permit to increase the number of protons for image reconstruction decreasing the total dose



THE ROLE OF MONTE CARLO

200 MeV, 179 projections at 1°, 5M Histories, 20 cm circular phantom

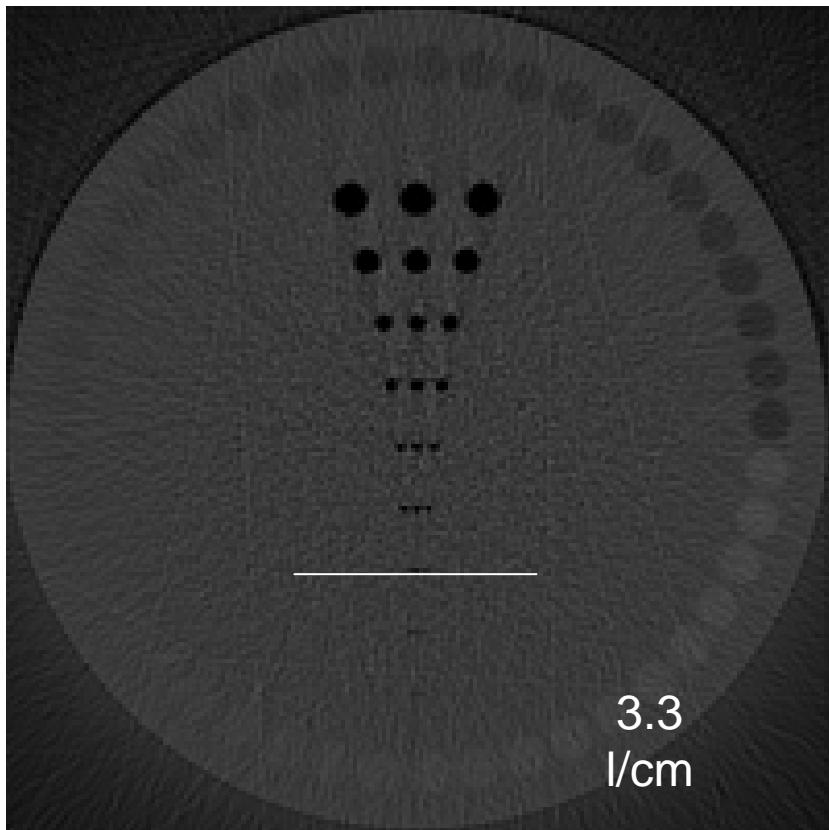
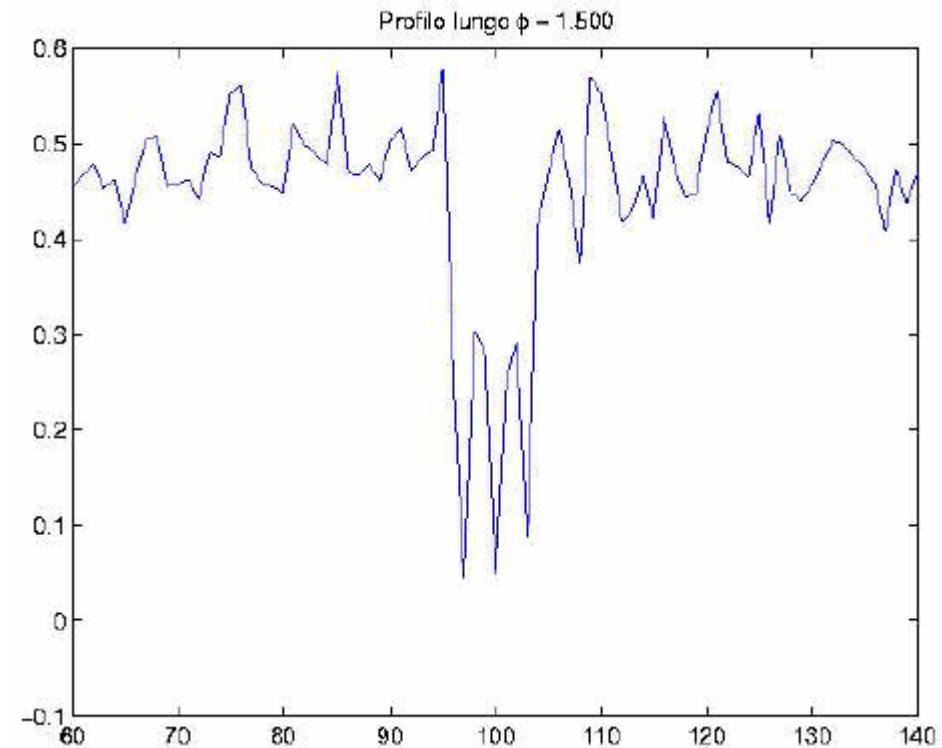
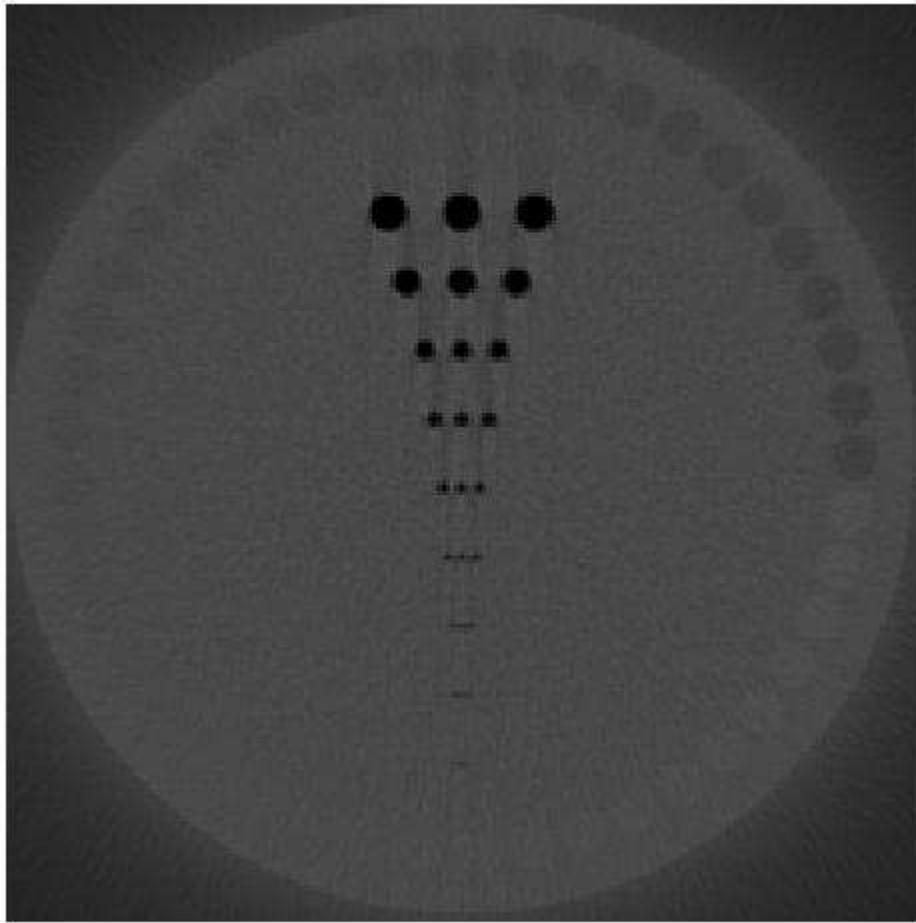


Image relative to
path1



THE ROLE OF MONTE CARLO

200 MeV, 179 projections at 1° , 250K Histories, 20 cm circular phantom



Maximum exit angle:
 1°

Image relative to
path2

THE ROLE OF MONTE CARLO

Table relative to image with path1

$N_p(x10^6)$	2.5	5	7.5
Dose(cGy)	15.5	31.08	46.62
$R_A(l/cm)$	3.3	3.3	3.3
$R_B(\Delta\rho/\rho_{H2O})$	4%	2%	2%

Table relative to image with path2

2.5×10^5

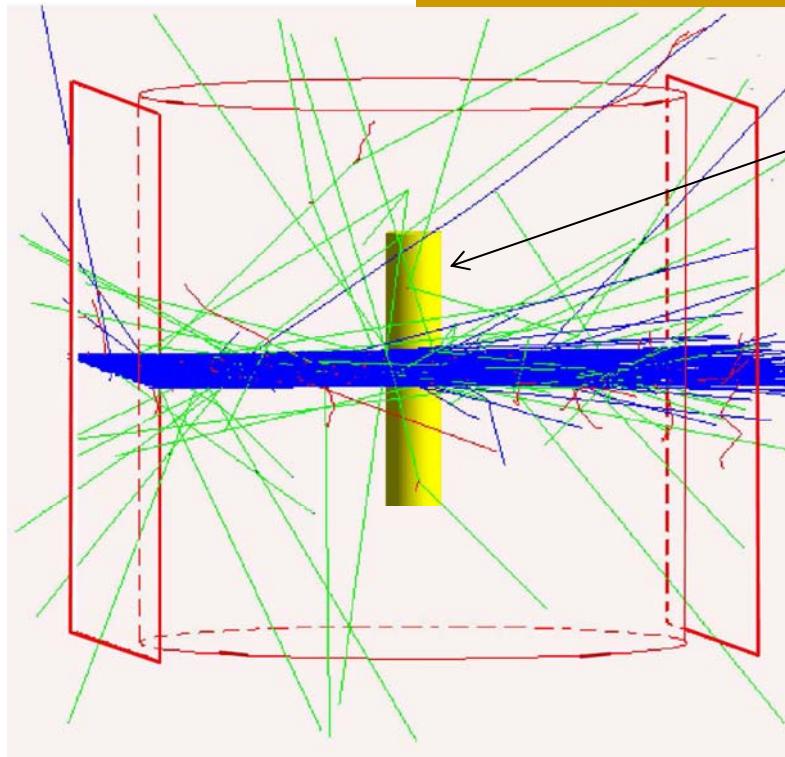
Dose (cGy)	5.1
$R_A (l/cm)$	2.5
R_B	4.5%

R. Shulte et al. *Reconstruction for proton computed tomography by tracing proton trajectories: A Monte Carlo study* Med. Phys. 33, 699 (2006).



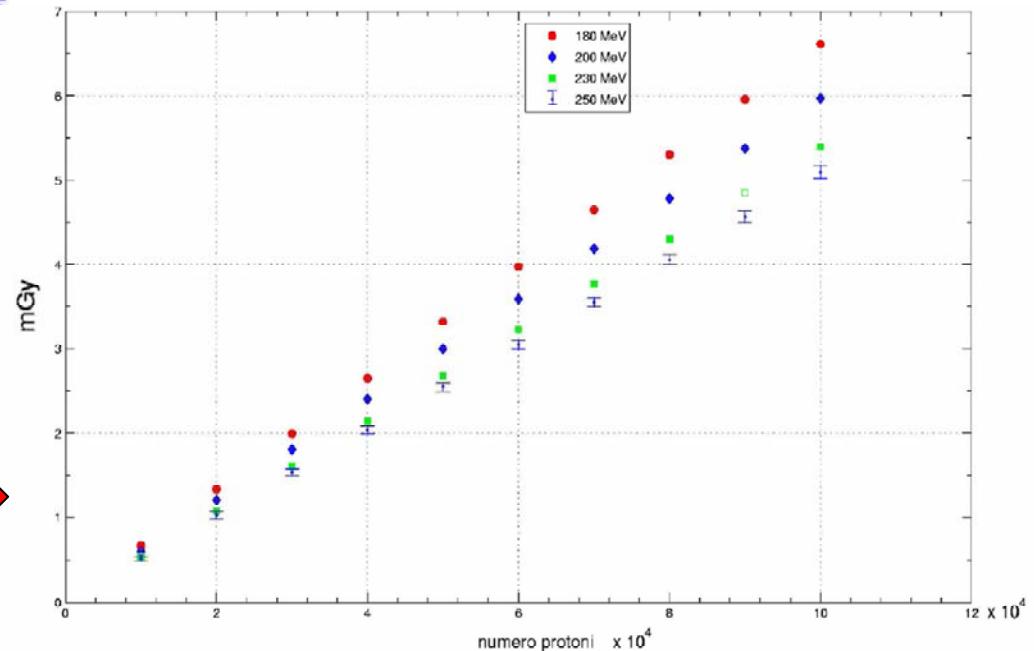
3-4 l/cm

Dose estimation: calculation of CTDI



Ionisation chamber

Dose versus number of histories

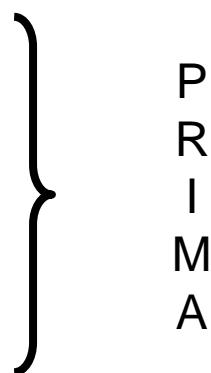


- ✓ Patient positioning
- ✓ Improvement of input information of TPS and, consequently, improvement in dose control
- ✓ Lower dose level as respect the conventional xCT

Good density resolution and very small dose levels

Not so good spatial resolution → improvement of path estimation and of reconstruction algorithms (ART)

ISTITUTION INVOLVED IN THE pCT

- Reinhard Shulte group at the Loma Linda University Medical Center
 - Eros Pedroni group at Paul Sherrer Institute (actually they stopped the program)
 - The Italian group author of the work I am presenting:
 - Laboratori Nazionali del Sud – INFN, Catania
 - Physics Department e INFN Section, Catania
 - Clinic Phisiophatology Department, Florence University
 - Energetic Department, Florence University
- 

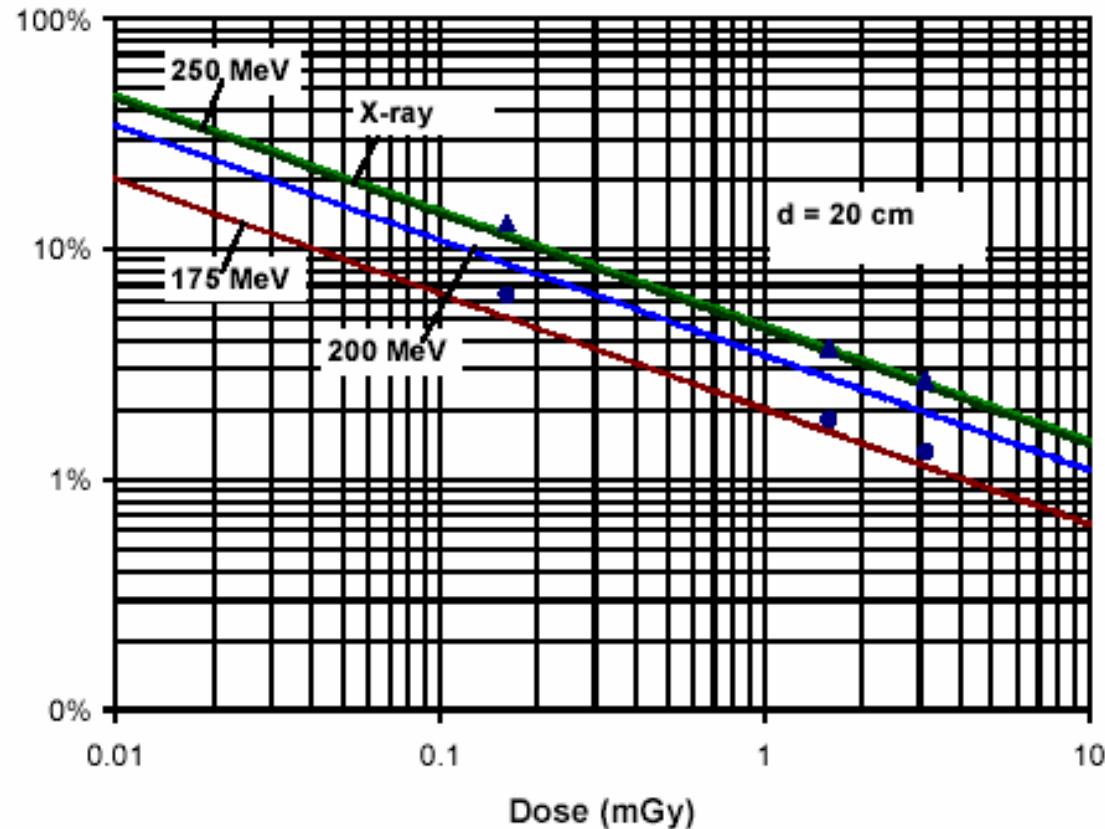


Thank you for your attention

LA RISOLUZIONE IN DENSITA'

Si richiede una accuratezza della conoscenza della densità elettronica in ogni voxel minore o uguale all'1%

Relative electron density resolution



GEANT4 studies:

R. Shulte et al.

Density resolution of proton computed tomography

Med. Phys. 32 (4), April 2005

PER ENERGIE OPPORTUNE E' POSSIBILE OTTENERE RISOLUZIONI IN DENSITA' EQUIVALENTI ALLA X-CT MA CON UN RISPRMIO DI DOSE