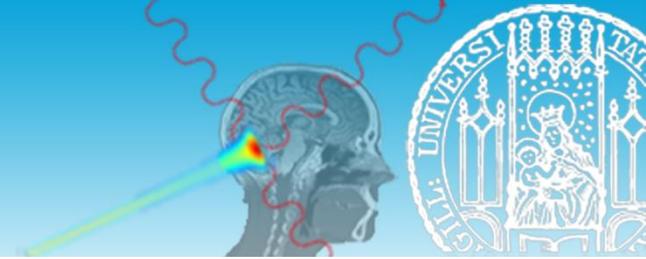


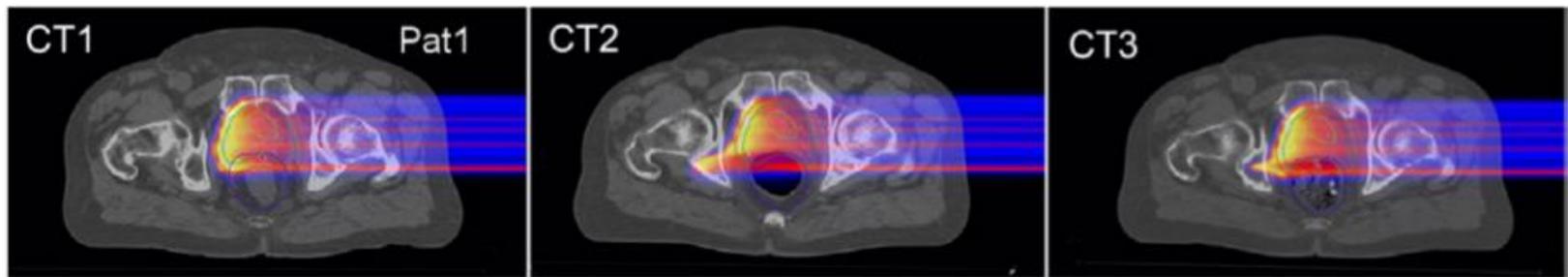


Toward a new Treatment Planning Approach accounting for in-vivo proton range verification

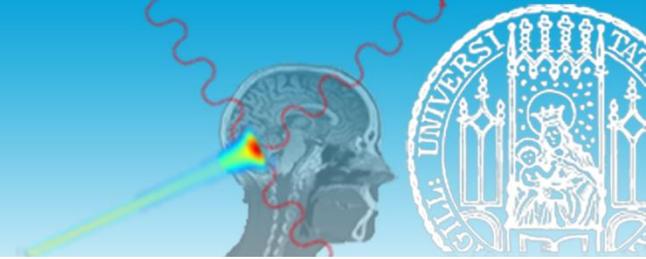
Ludwig-Maximilians-Universität, München
Liheng Tian



- Proton range uncertainties



- CT number conversion
- Anatomical changes
- Patient set up
- ...

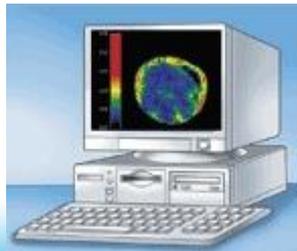
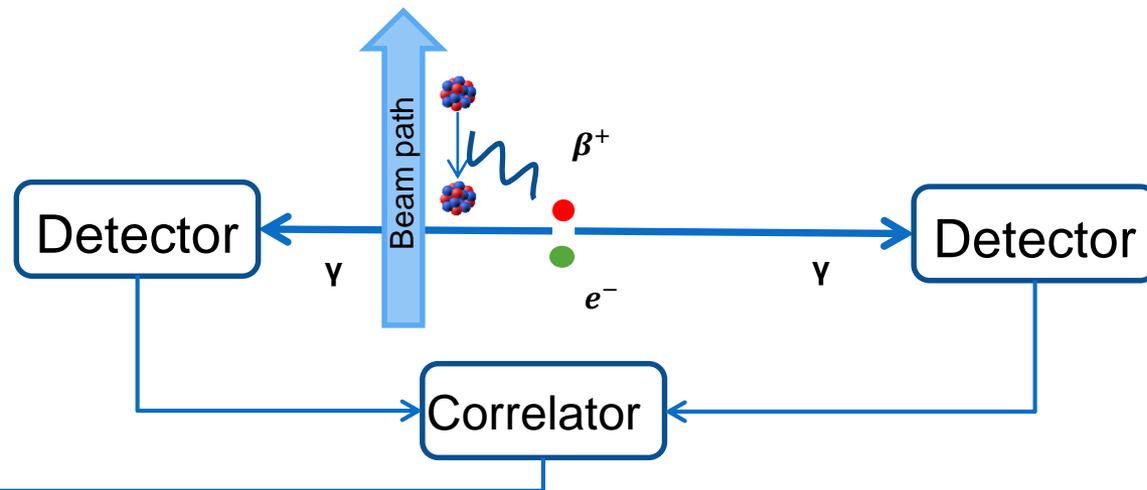


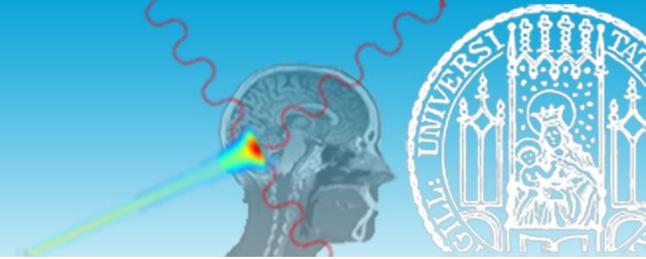
• In-vivo proton range verification

• Positron Emission Tomography (PET)

Detection of annihilation photons from the decay of β^+ -emitters

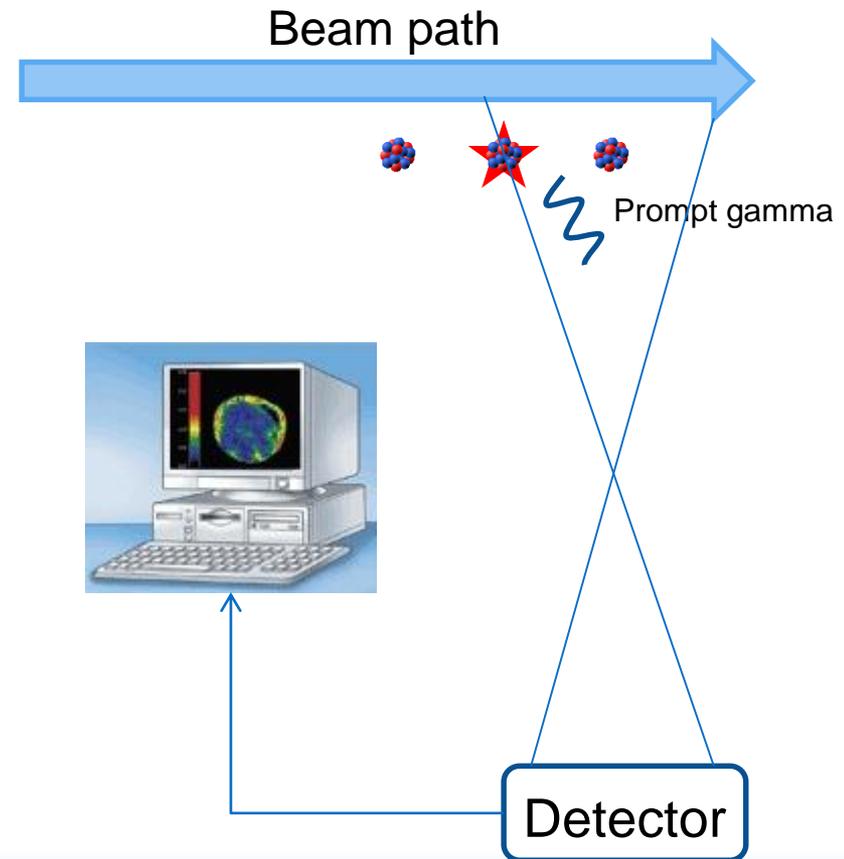
1. Takes few tens minutes to use PET scanner
2. Disturbed by organ motion and washout effect



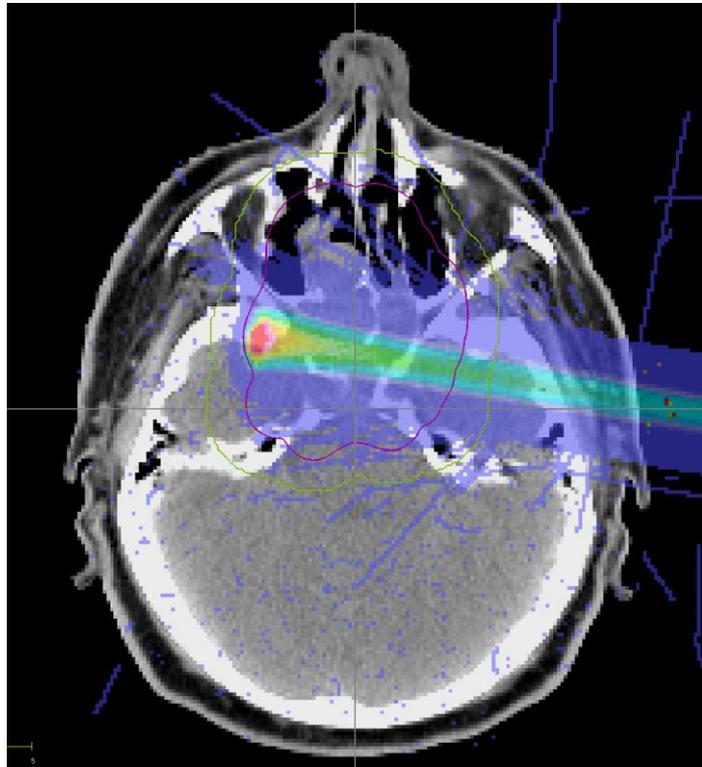
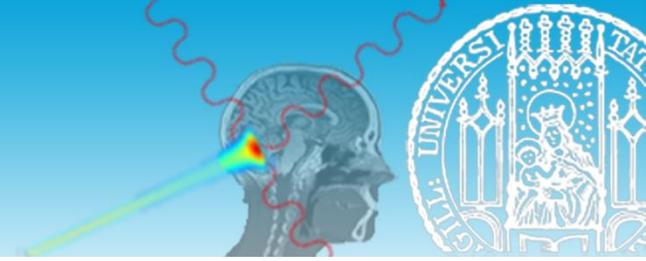


• In-vivo proton range verification

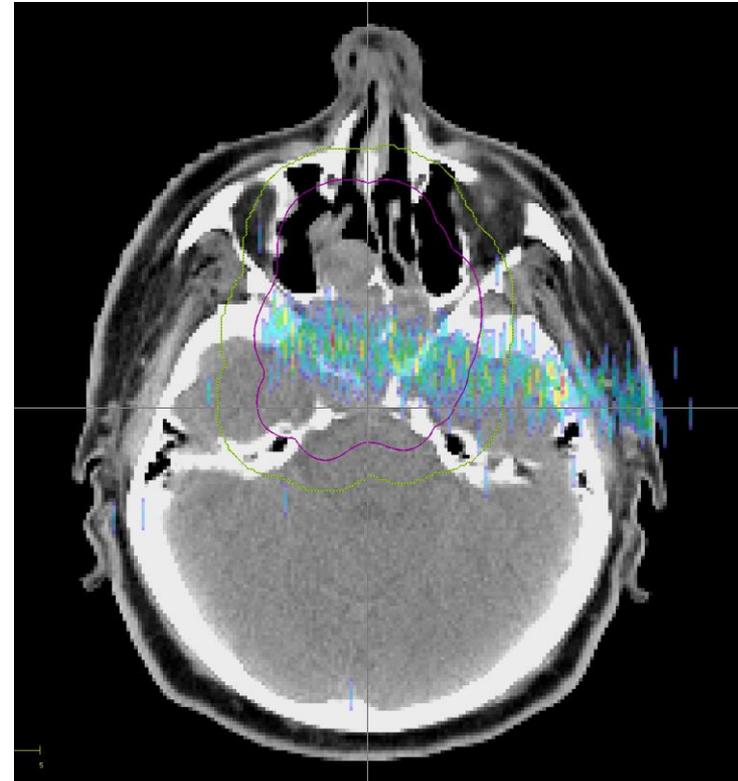
- Prompt Gamma (PG) Imaging
 - Monitor the Prompt Gamma emitted by nuclear de-excitation of nuclei in the beam path
 - 1. Fast (within nanoseconds)
 - 2. Better PG-dose signal confirmation
 - 3. Higher gamma production rates
 - 4. Clinical experiment shows that the retrieval precision of 2mm can be achieved



Introduction

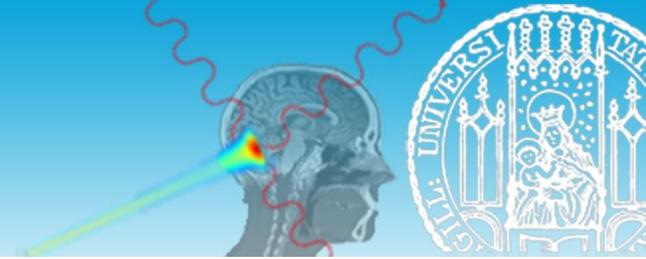


Dose delivery

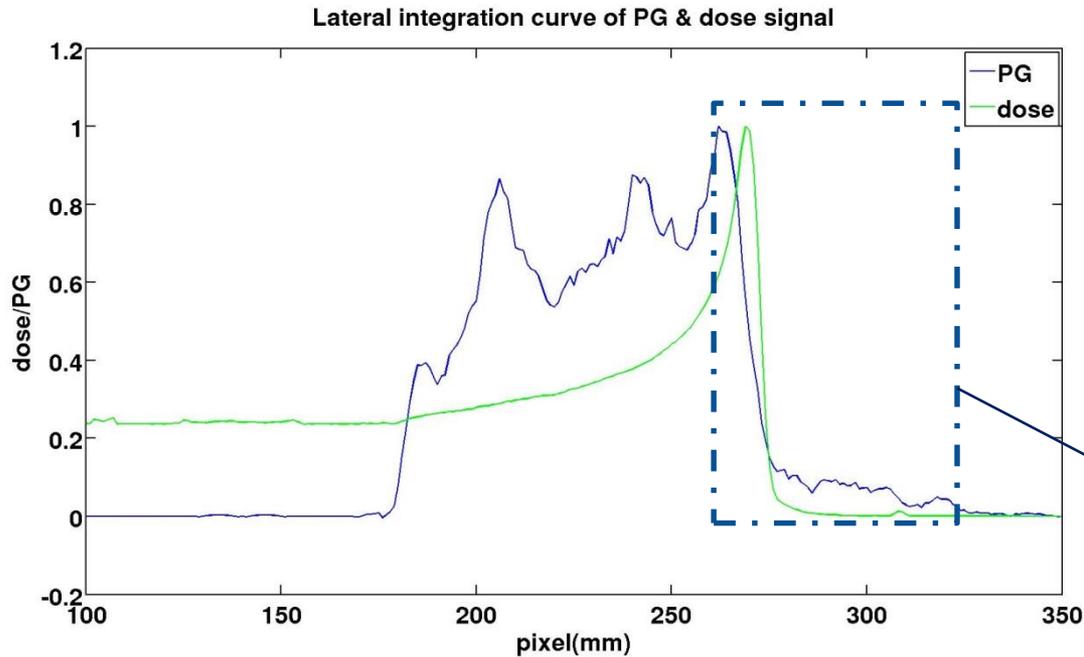


PG emission

Introduction



- Ideal PG & dose profile

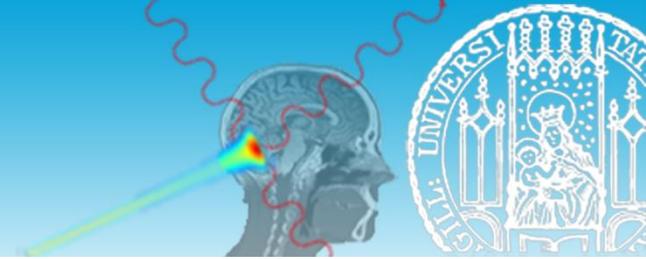


Lateral integration of PG & dose signal

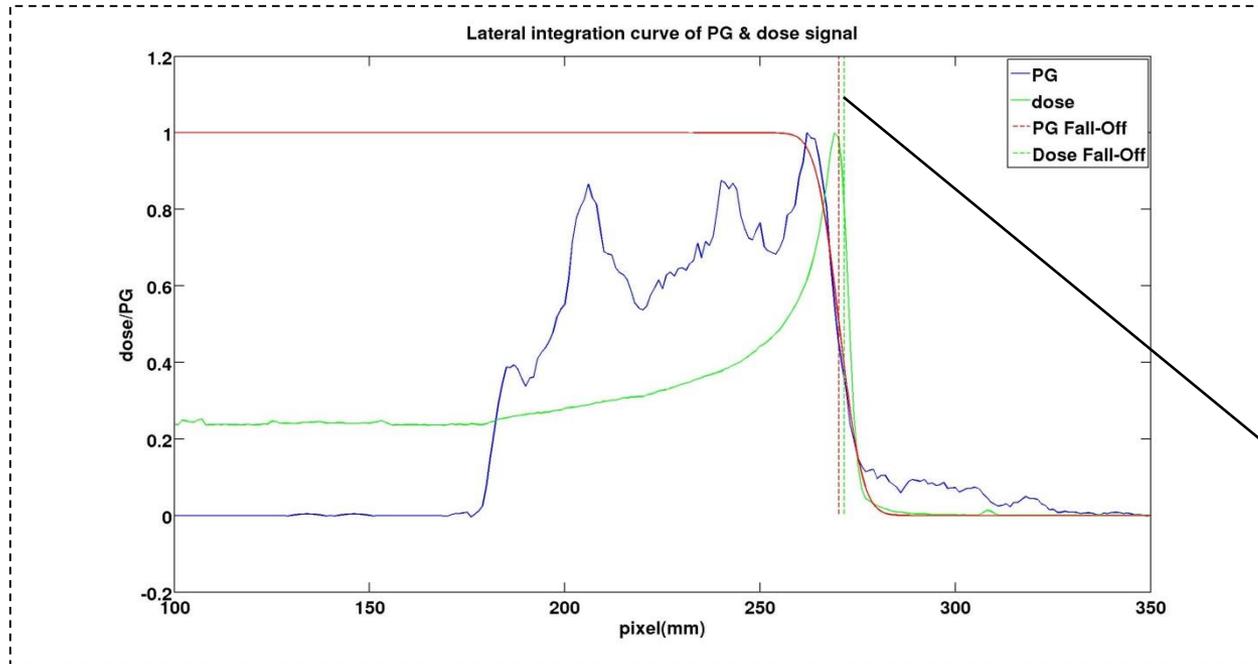
c: Fall-off position
1/b : Fall-off slope

$$F(x) = a + (1 - a)\text{erf}[b(x - c)]$$

Introduction



- Ideal PG & dose profile

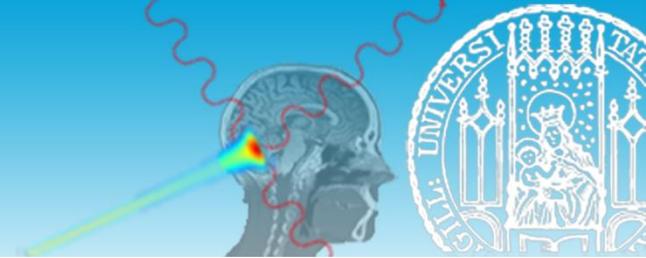


$$dR = -1.30 \text{ mm}$$

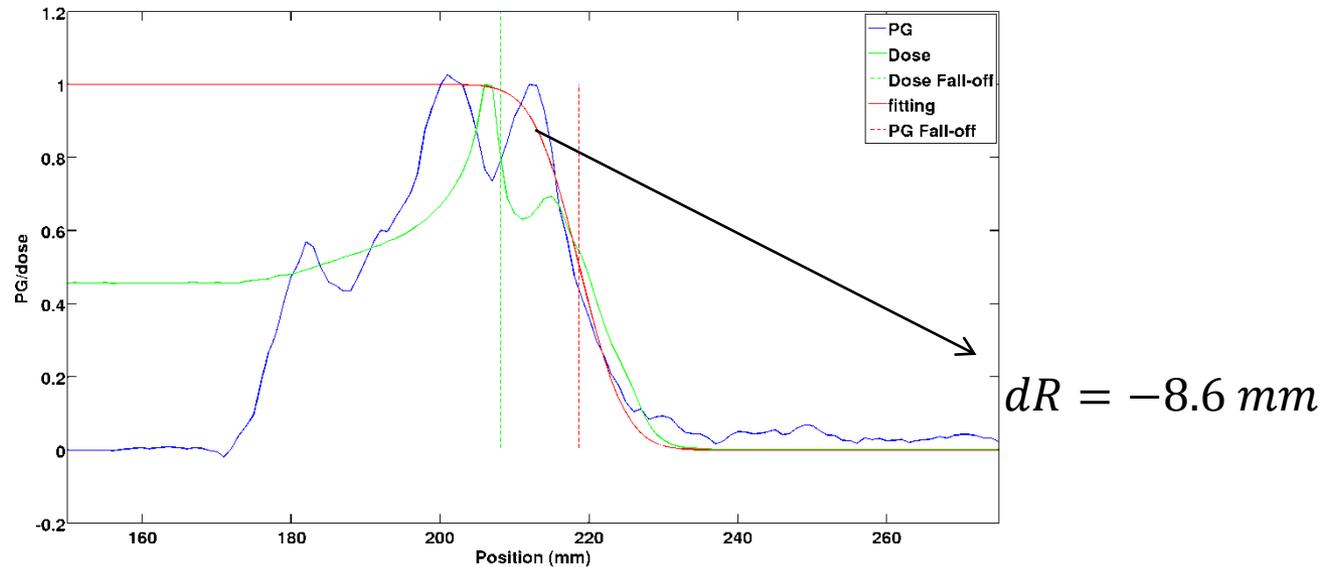
Lateral integration of PG & dose signal

Distance between PG and dose fall-off position:
 dose: distal 80% dose fall off
 PG : fitting function

Introduction

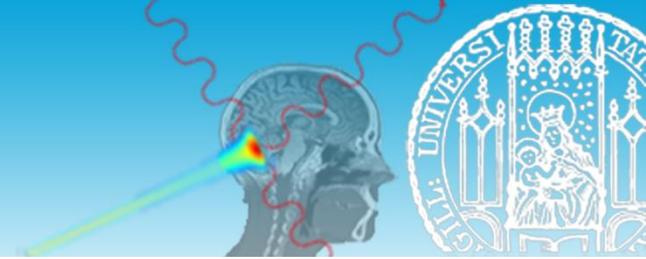


- However, the correlation between PG and dose signal is not always good



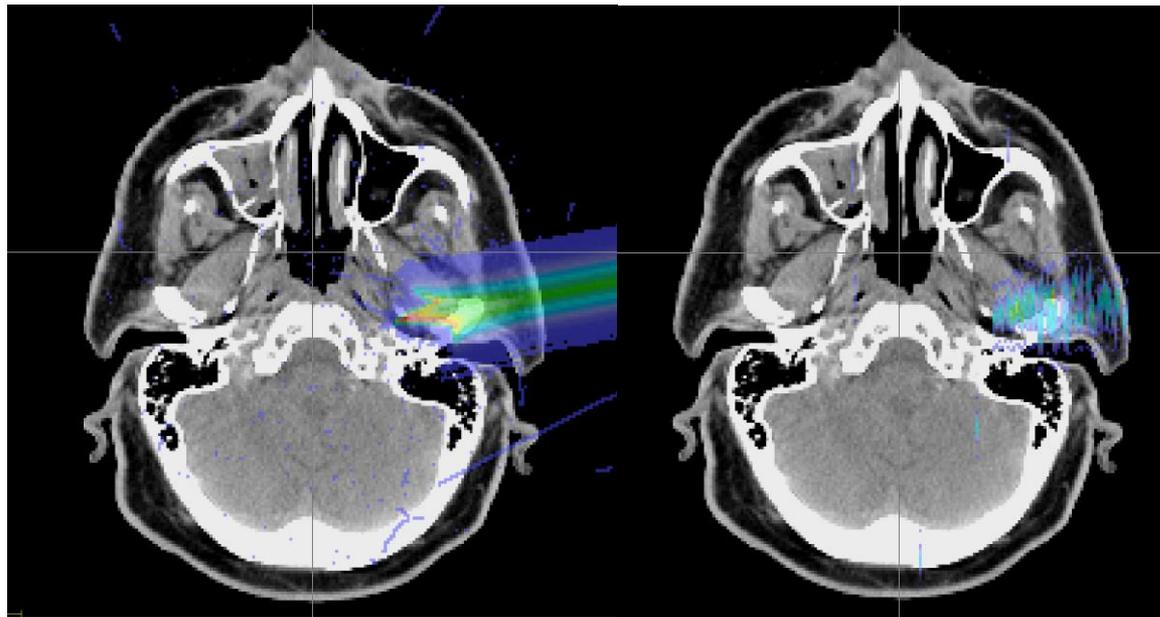
Lateral integration of PG & dose signal

Introduction



- Reason

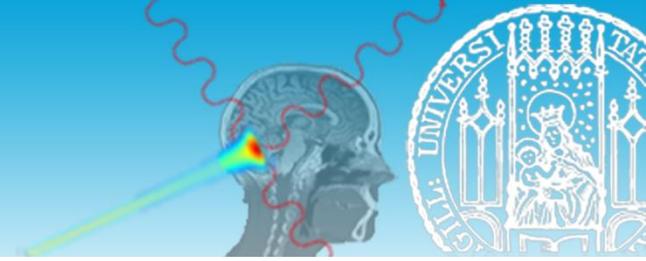
Tissue inhomogeneity (bone/air in beam path)



Dose delivery

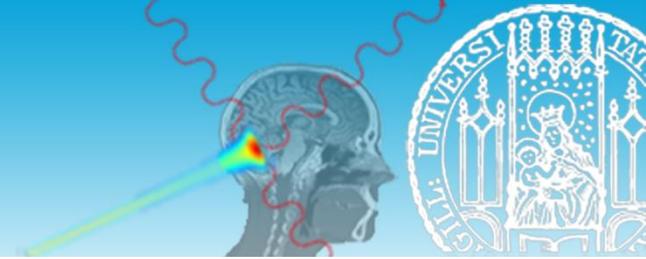
PG emission

Introduction

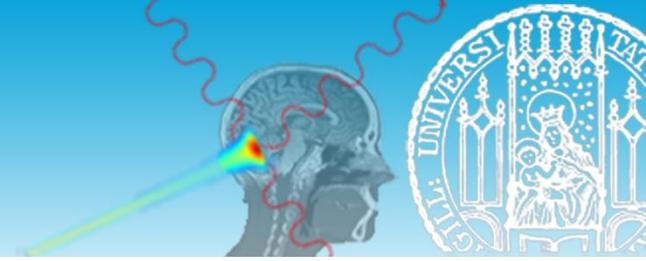


- The PG-dose correlation of individual pencil beam is not considered in current treatment planning system (TPS)

Goal



- Goal of my study:
Optimize current TPS system, accounting for in-vivo proton range verification to enable a reliable treatment monitoring

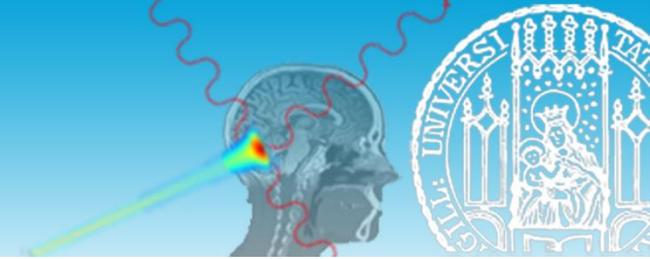


1. Quantify and identify the degree of 'PG-dose correlation' for each spot

Criteria:

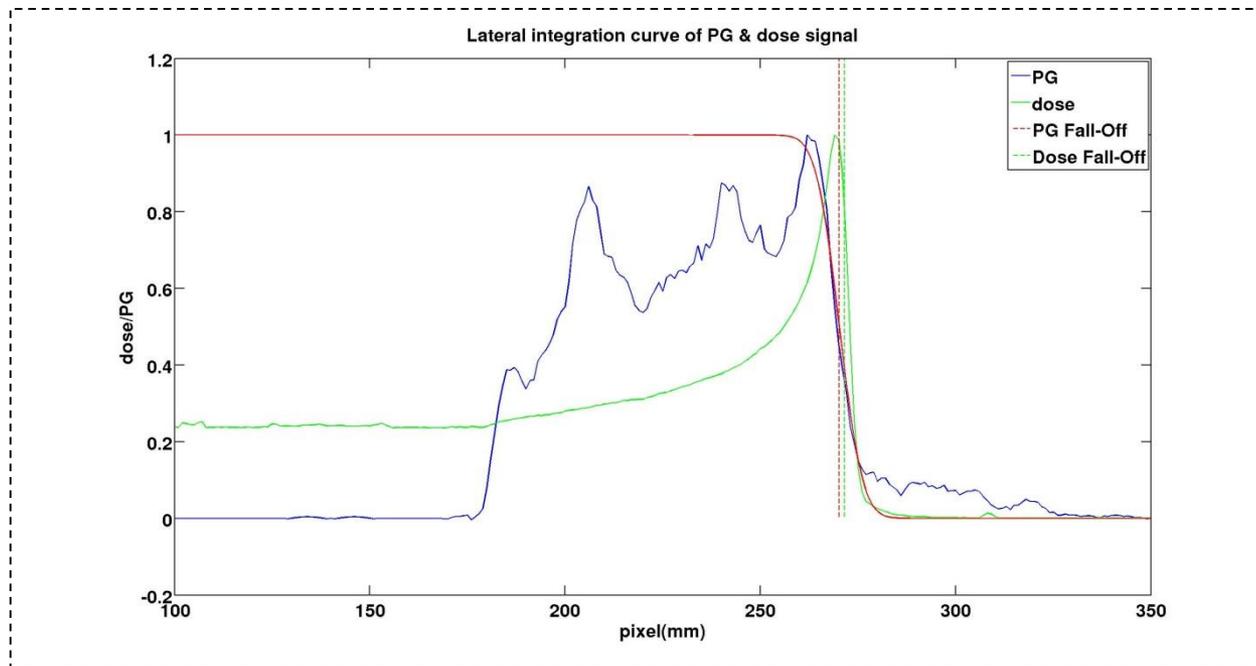
- Distance between the Fall-off positions of PG & dose profile (dR)
- Slope of the PG signal Fall-off region ($1/b$)
- Fitting quality (Sum of squared errors, SSE)

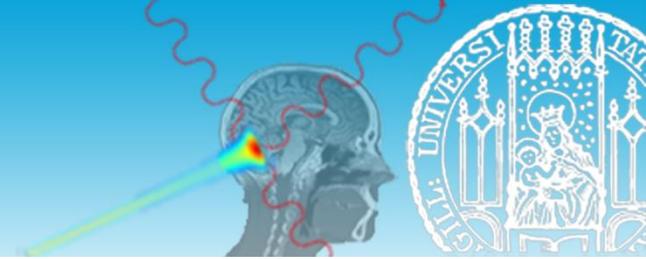
Method



- Example:

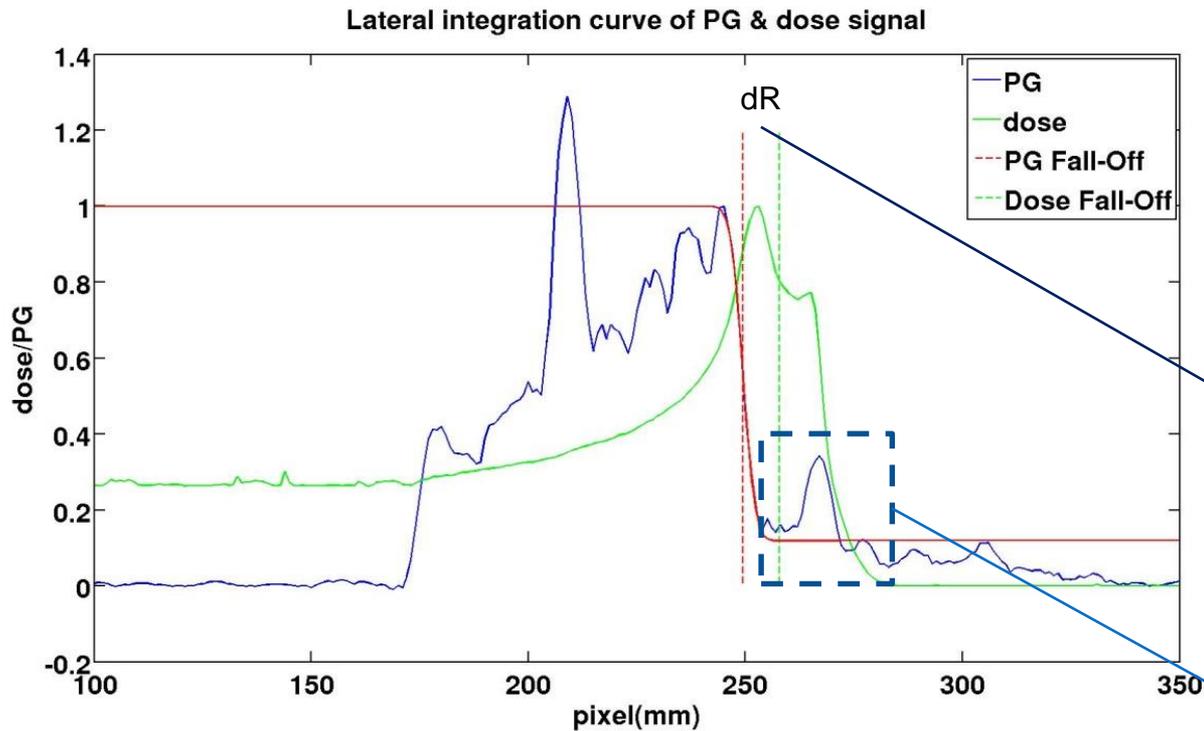
Ideal case:





- Example:

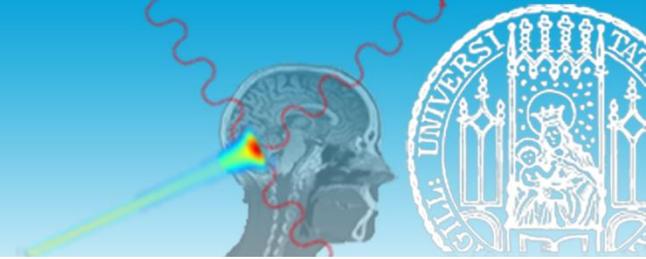
Poor correlation case 1:



Large dR relates to poor correlation directly

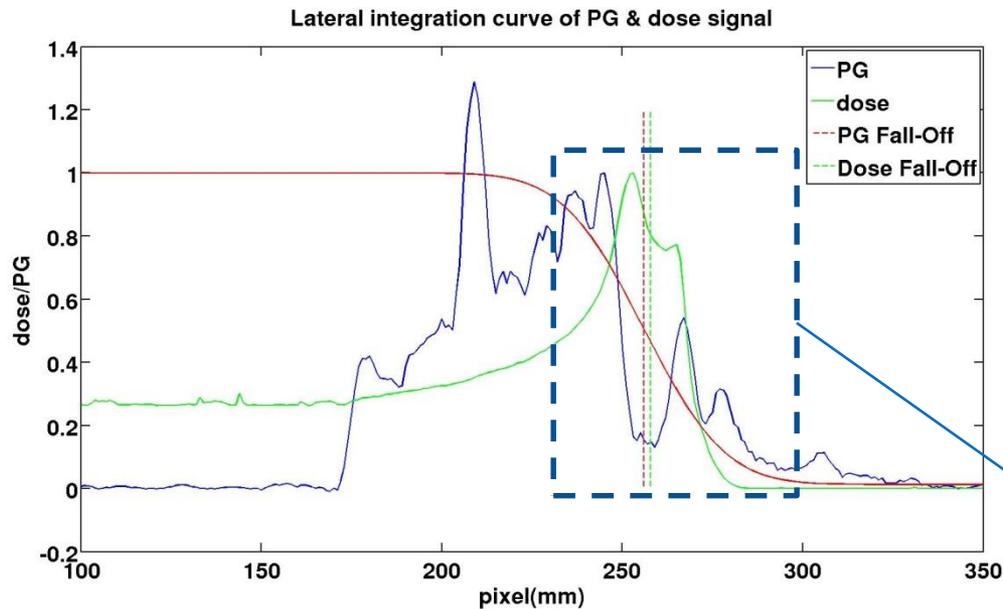
A secondary peak, relating to an air cavity, leads to high SSE value

Method



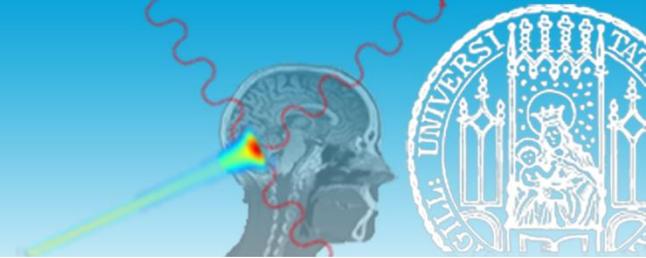
- Example:

Poor correlation case 2:

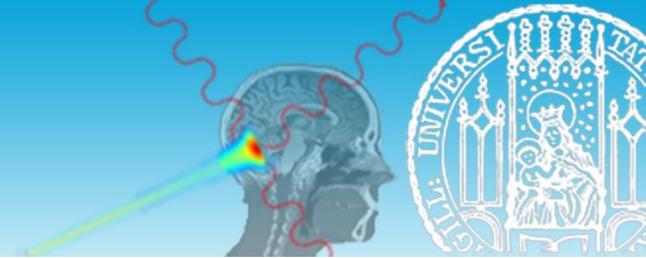


A shallow slope related also to an air cavity, causing the failure of fitting

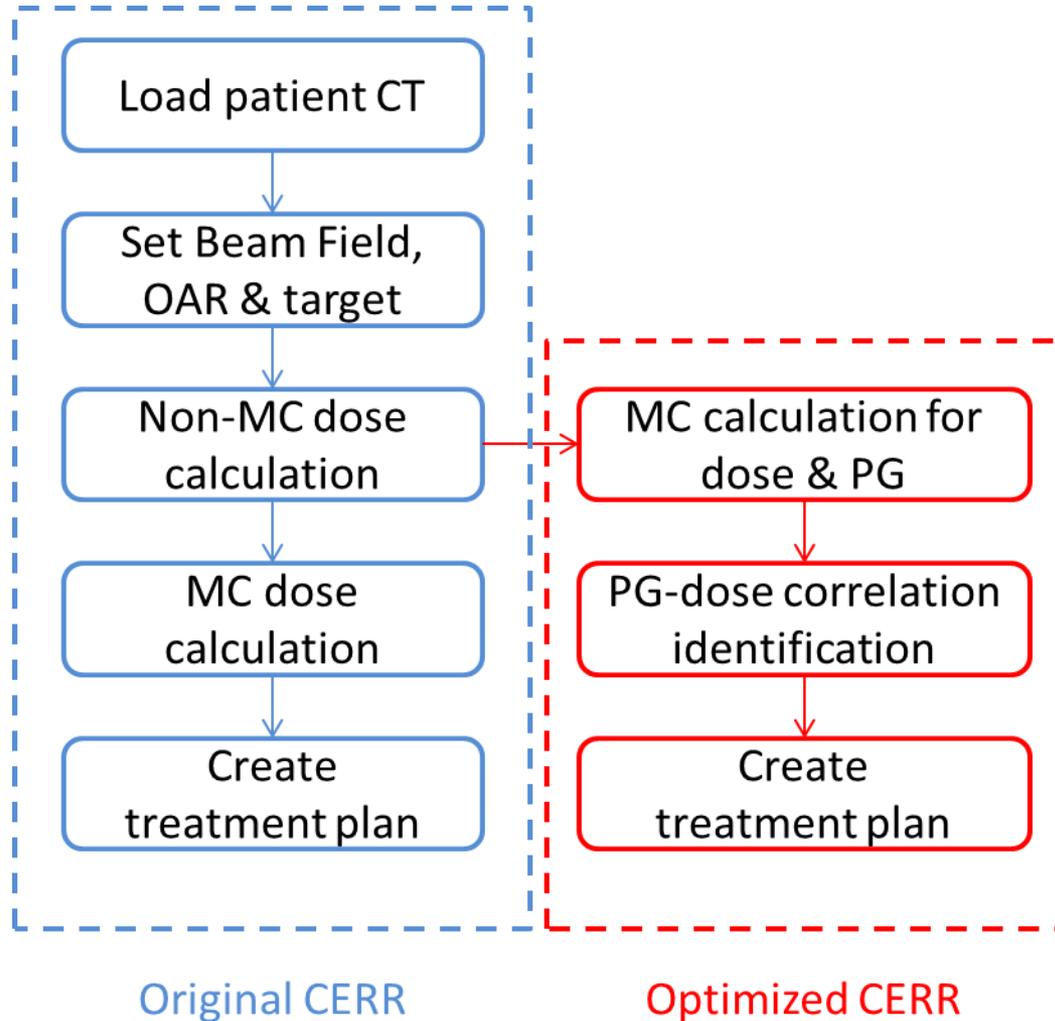
Method



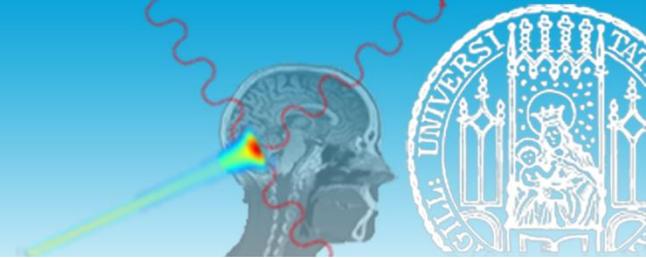
1. Quantify and identify the degree of 'PG-dose correlation' for each spot
2. Improve the initial treatment plan while maintaining the quality of plan



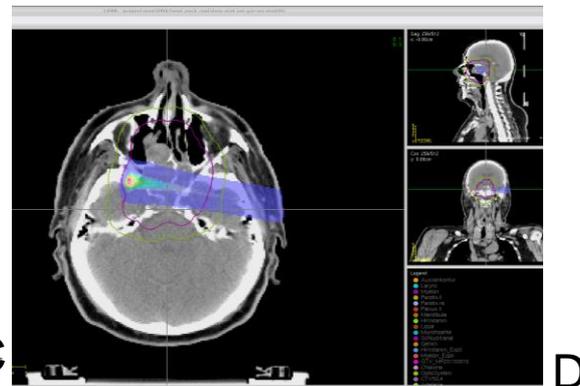
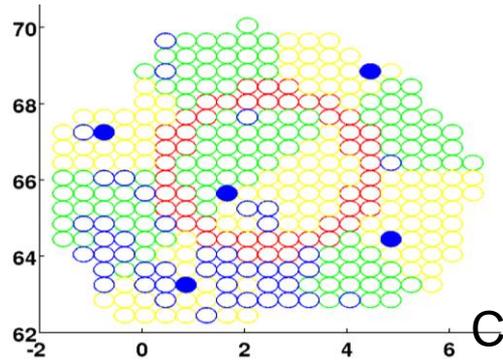
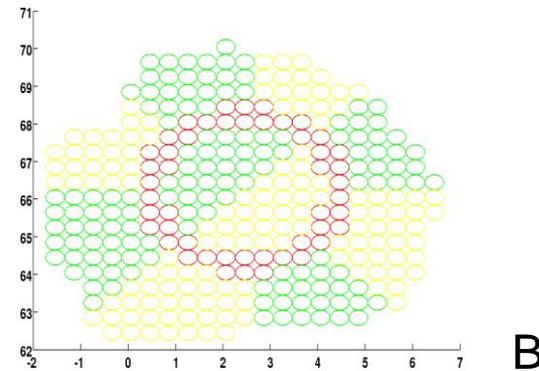
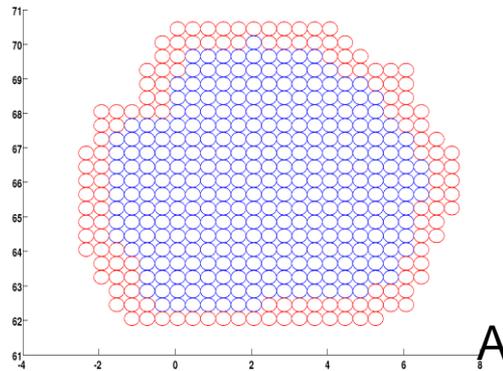
General workflow of CERR TPS and our improved TPS:



Method



- Boost few spots which have the best PG-dose correlation

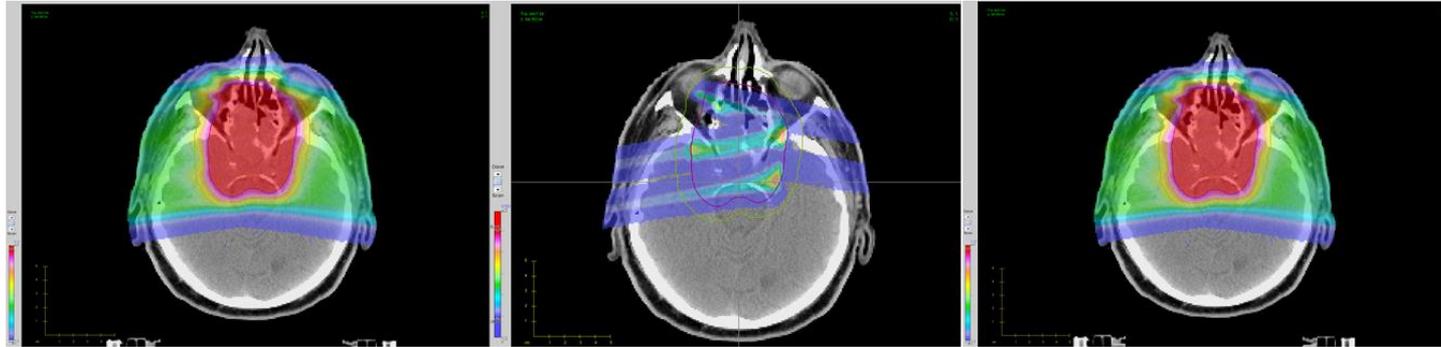


Result

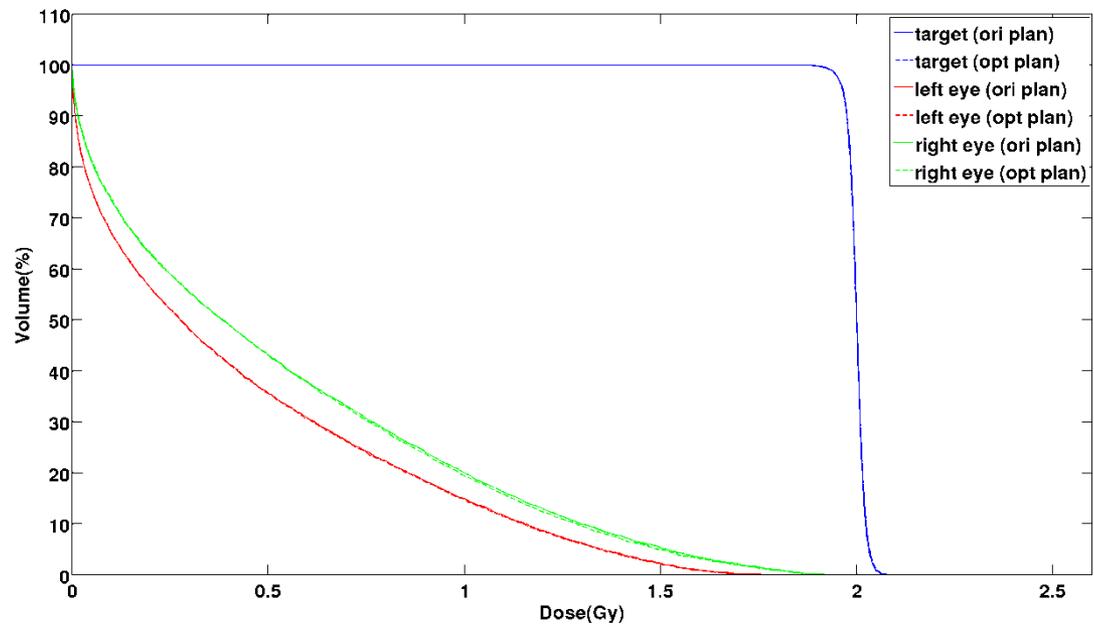


- Dose delivery
- PG-dose correlation

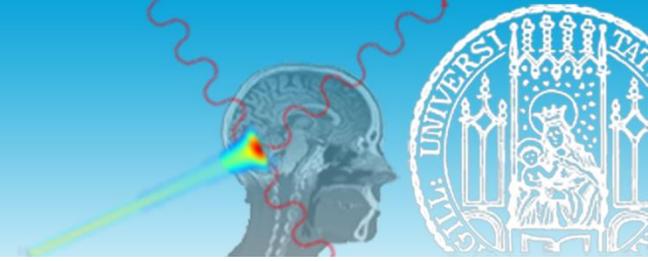
Dose delivery



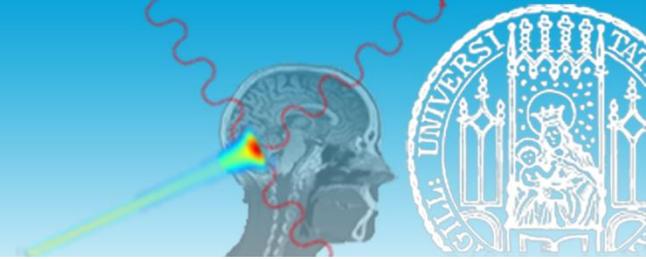
Patient
1



Result

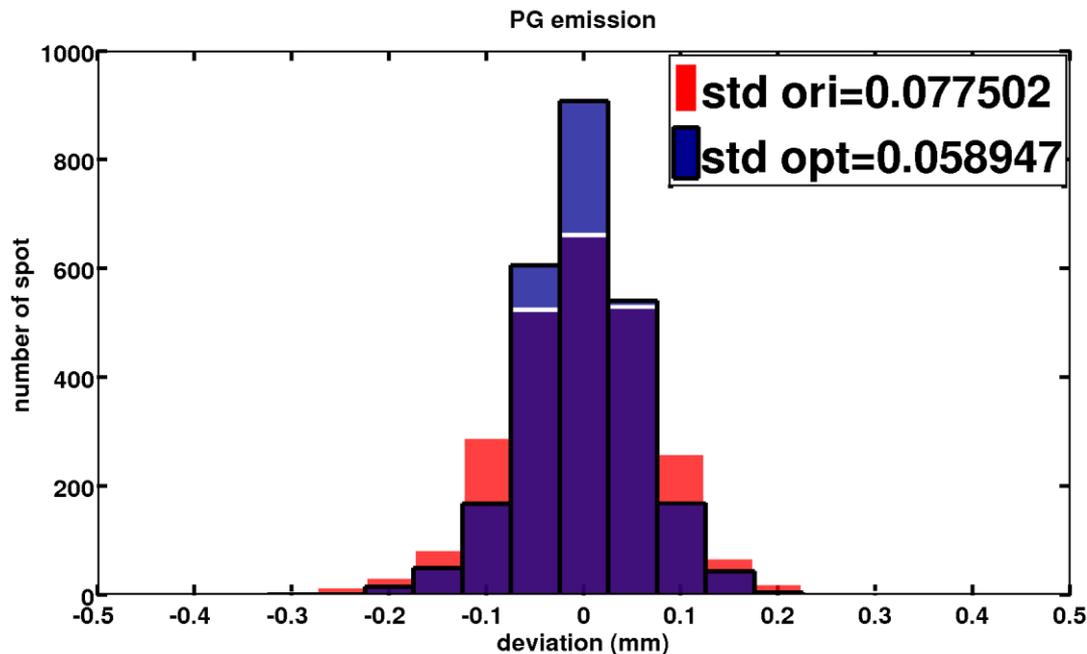


- Dose delivery
- PG-dose correlation

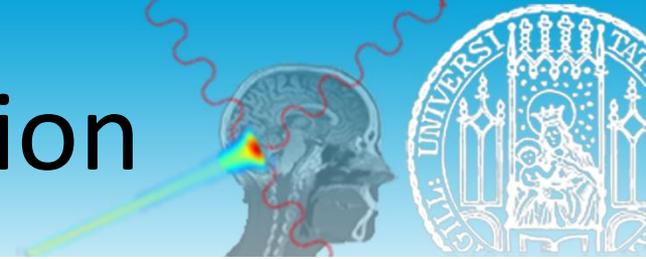


Robustness of fitting:

10 highest-intensity spots in both plans are simulated 25 times, the number of protons in each individual simulation is 10^7 . Fitting for each individual simulation (10^7) is compared to full simulation (10^8) of the same spot to evaluate the robustness of fitting.



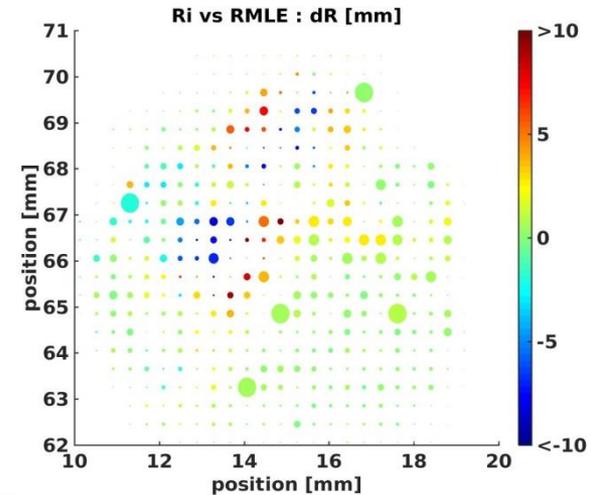
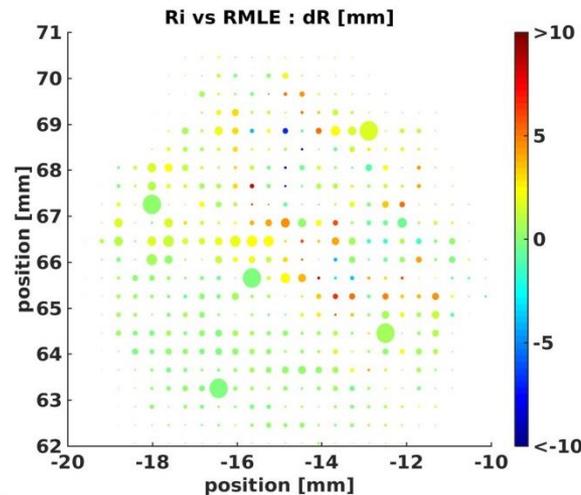
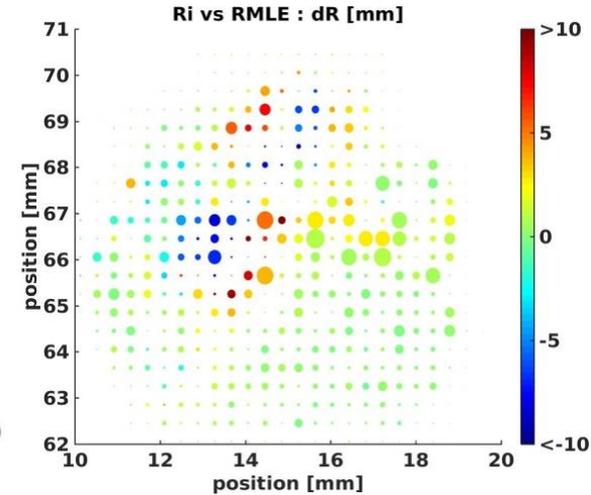
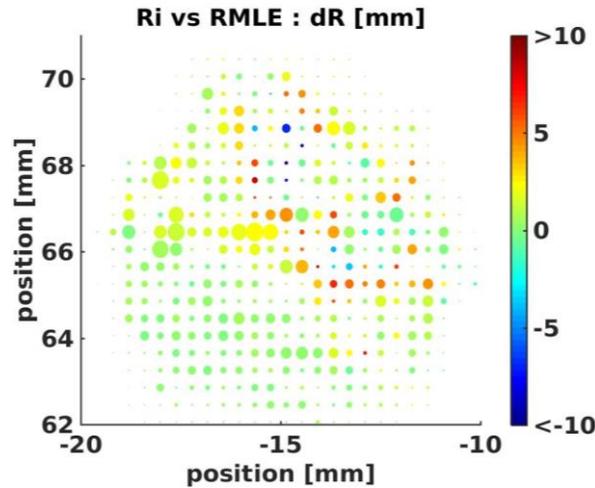
PG-dose correlation

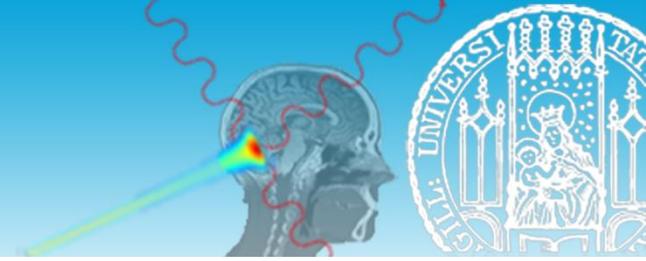


Beam's eye view:

The color shows
the PG-dose
correlation and

The spot radius is
linearly proportional
to the proton
number

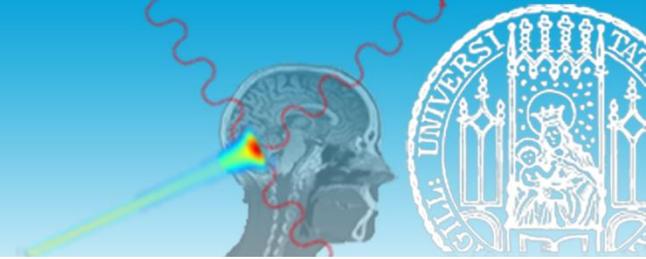




Currently, our new approach is able to:

- Quantify and identify spot-by-spot PG dose correlation automatically in a MC treatment planning process
- Provide the user with candidate spots which have reliable PG-dose correlation
- Optimize a treatment plan which is comparable to the initial treatment plan in terms of dose distribution. Besides, few high-intensity spots in the new plan could provide reliable in-vivo proton range verification

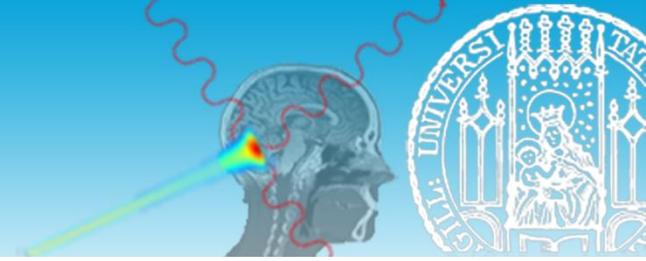
Conclusion



Though different issues are still needed to be addressed, with the results of this project, the treatment planning process in proton therapy could be improved, by integrating the PG-based in-vivo monitoring of the beam range

This novel approach could lead to a safer and more controllable proton treatment in the future

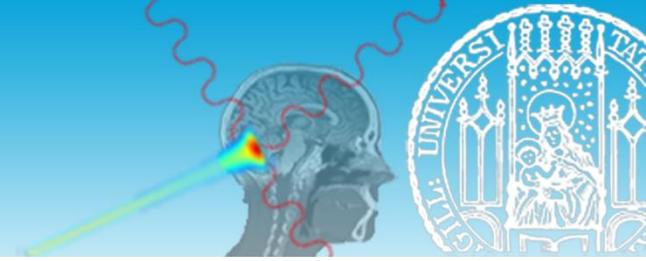
Current work



- Detection devices signal is different from PG emission
- Robustness of PG-dose correlation considering anatomical change and different uncertainties



LUDWIG-
MAXIMILIANS-
UNIVERSITÄT
MÜNCHEN



Thank you!