

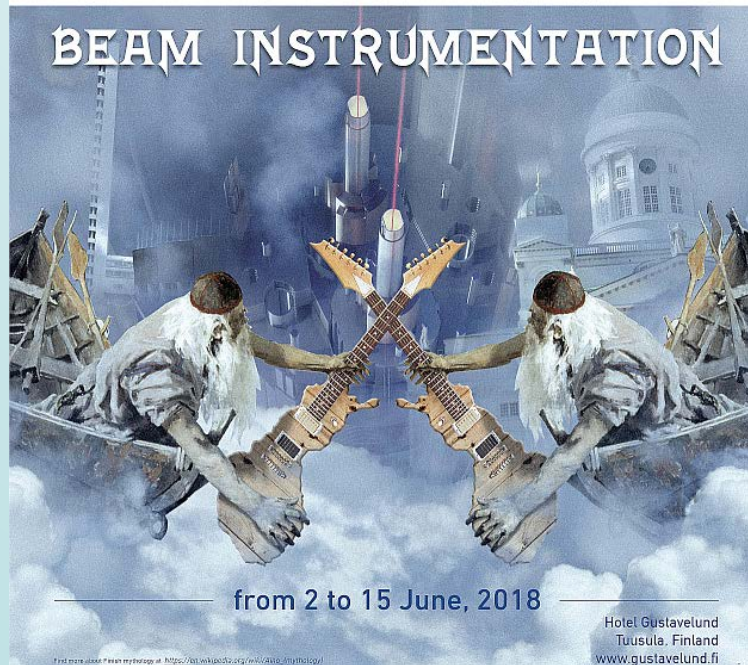
# Medical Applications Instrumentation & Diagnostics

*Andreas Peters*

*(HIT, Head Accelerator Operations)*



## BEAM INSTRUMENTATION



HIT Betriebs GmbH am  
Universitätsklinikum Heidelberg

<http://www.hit-centrum.de>

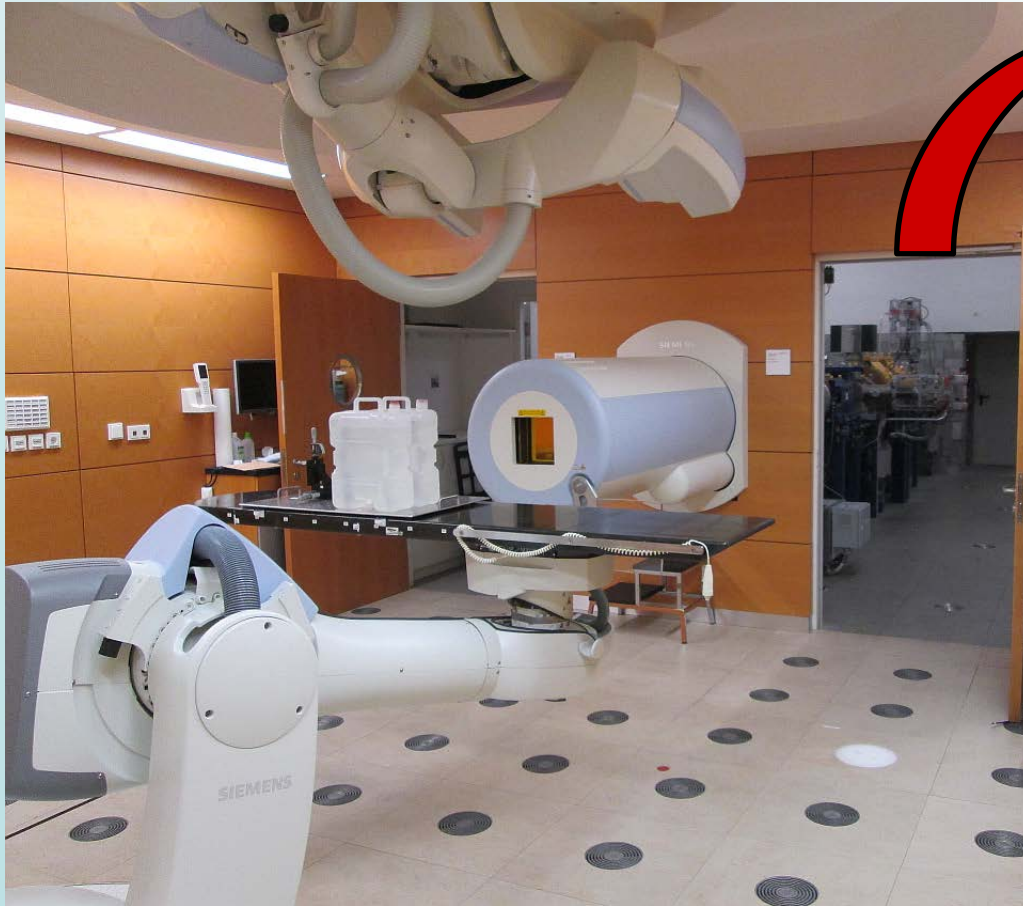
# Outline

- The role of beam instrumentation and diagnostics in particle therapy accelerators
- An extended view: Instrumentation – Feedbacks – Detector Technology – Quality Assurance
- Diagnostics and instrumentation techniques used in particle therapy: the interaction of accelerator and irradiation technology – some basics, examples and challenges in near future
- Conclusion and Acknowledgements

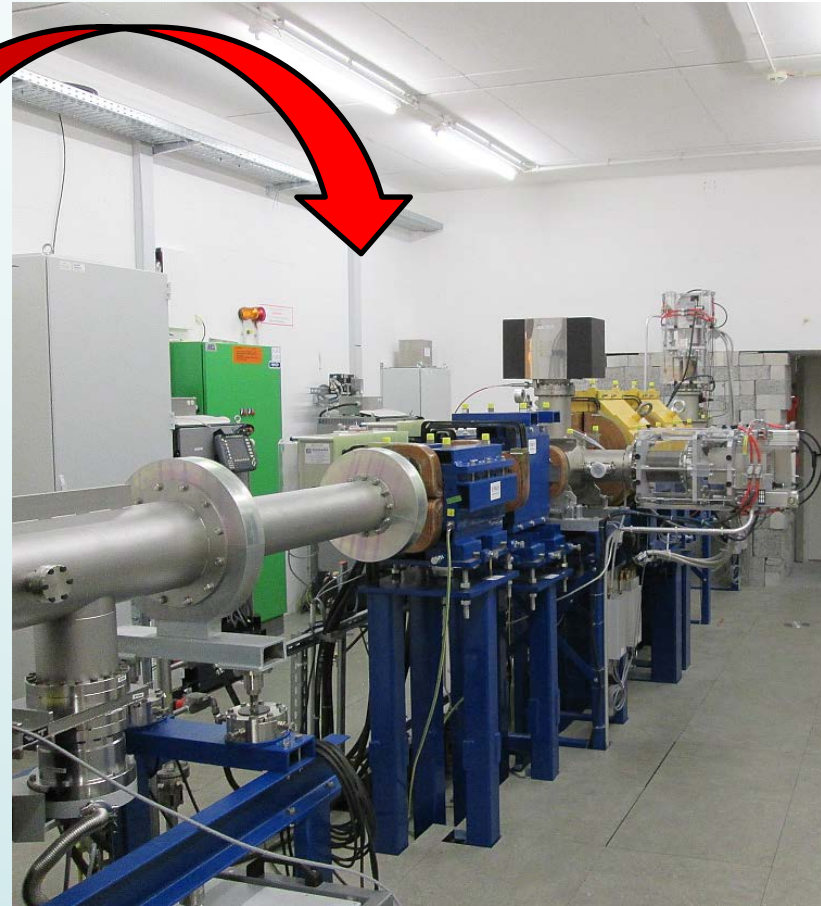
# Beam instrumentation and diagnostics in particle therapy accelerators

- Is there any difference between a scientifically used accelerator and a medical accelerator? **No, not at all!**
- Due to the economic effects sometimes only the absolute **minimum of beam diagnostics** is installed for commissioning and **standard operation**.
- Most important devices are **current transformers** for **online diagnostics** (non-destructive) and **beam profile monitors** for the **daily machine QA** (destructive) – these topics are covered by other talks of this CAS.
- The **speciality of medical accelerators** is the **strong link to the beam consumer**, in this case the **irradiation technology** and the high-conformal **dose application**.

# Beam instrumentation and diagnostics in particle therapy accelerators



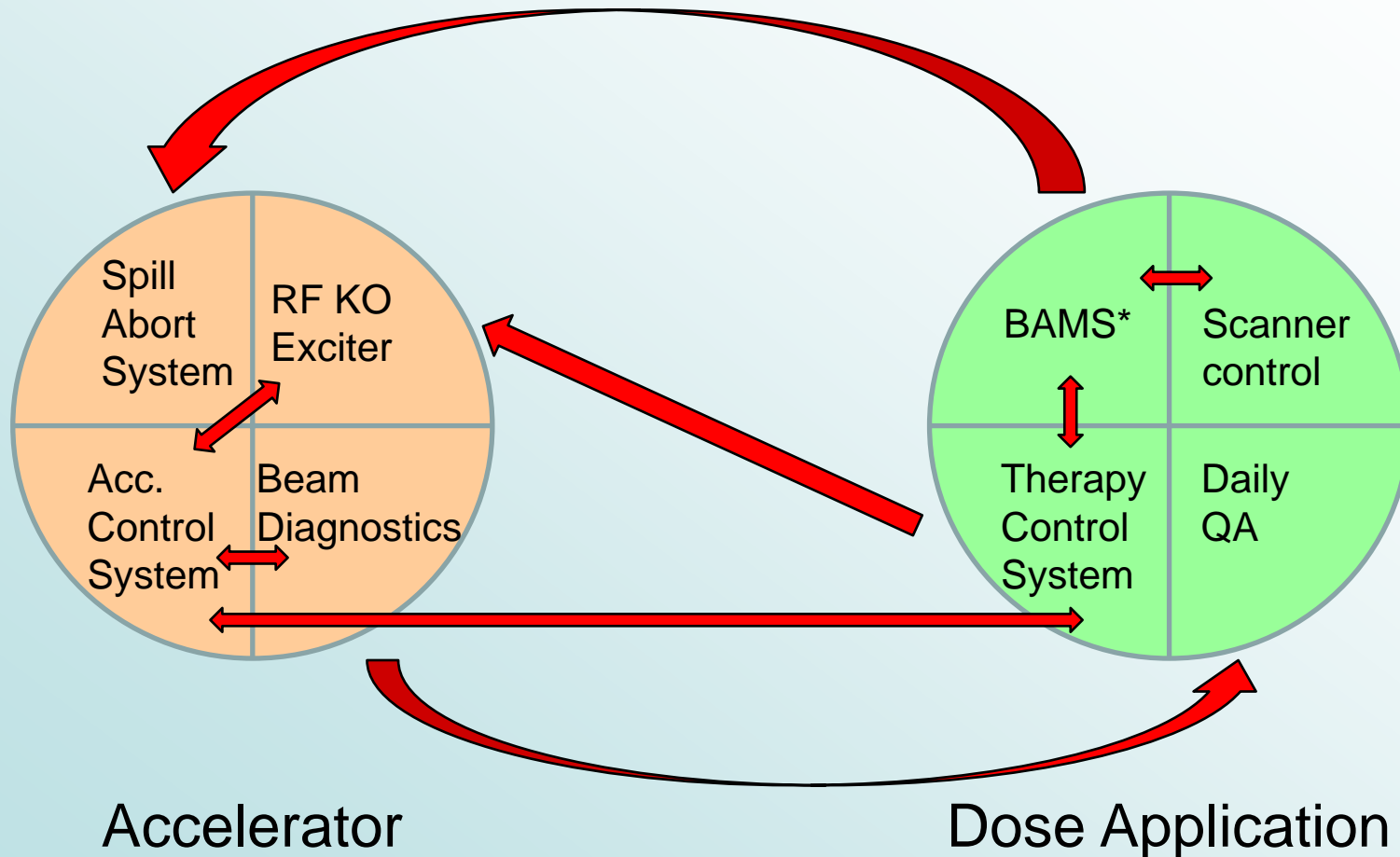
Horizontal patient treatment room at HIT



Beamline with quadrupoles and scanner magnets behind

# An extended view

- Instrumentation – Feedbacks – Detector Technology – Quality Assurance



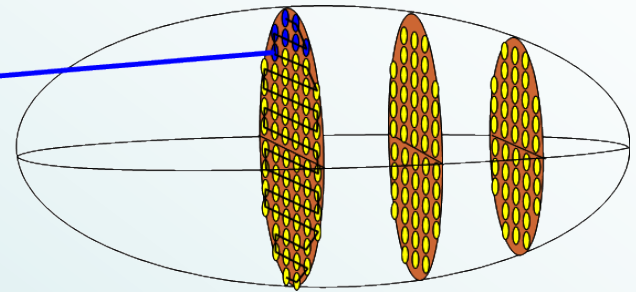
\* Beam Application and Monitoring System

# Interaction of accelerator and irradiation technology in particle therapy – some basics

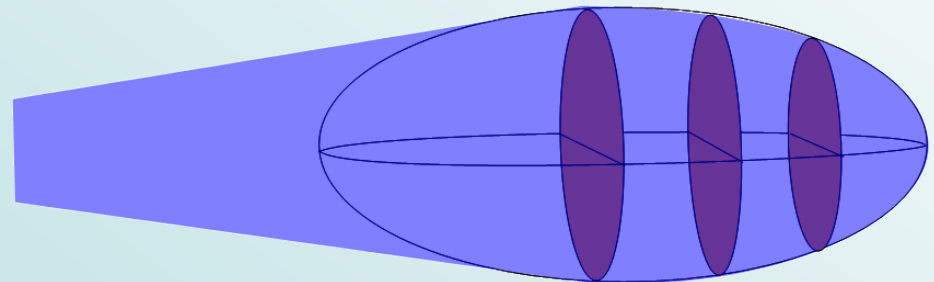
# Raster scanning technique

(also named *pencil-beam scanning*)

“Scanning” in the context of particle therapy is the application of the desired dose by means of a **pencil beam**, which is **moved** over the target point-by-point to cover the whole target volume.

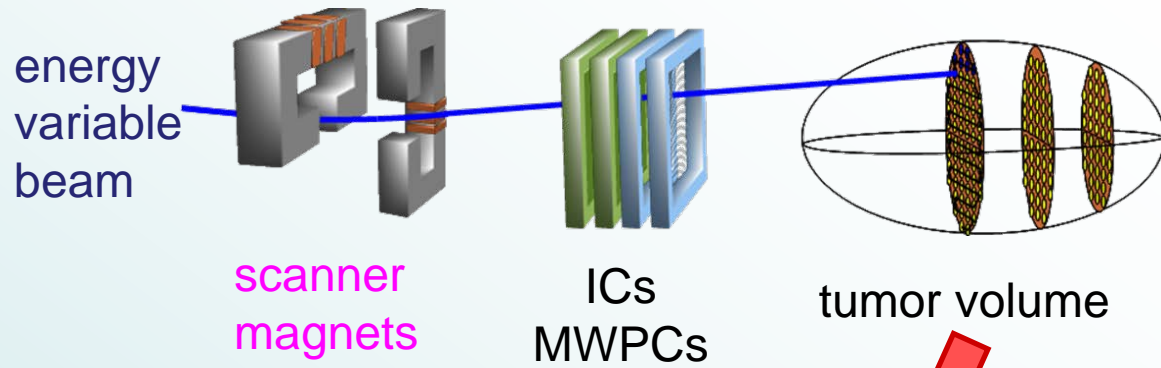


This is in contrast to broad-beam-techniques that use an expanded particle beam to irradiate the whole area or volume of the target at the same time.



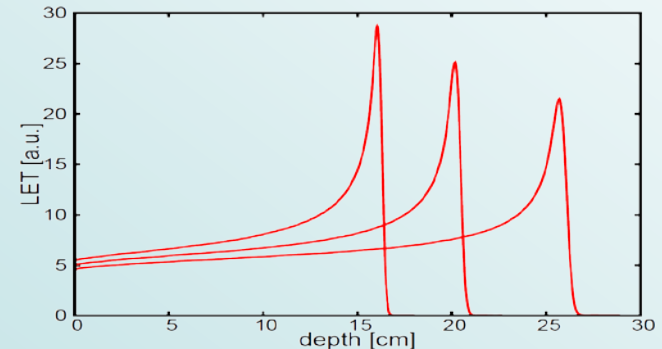
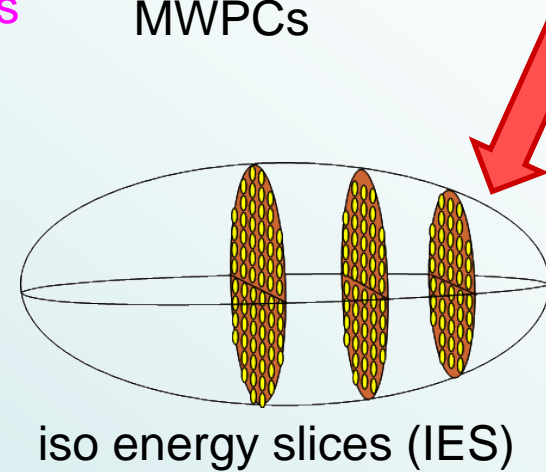
# Raster scanning technique

## Raster scanning system



The part of a treatment plan corresponding to a fixed (water equivalent) depth is called an **iso-energy-slice (IES)**.

Selection of iso energy slices is achieved by varying the beam energy.

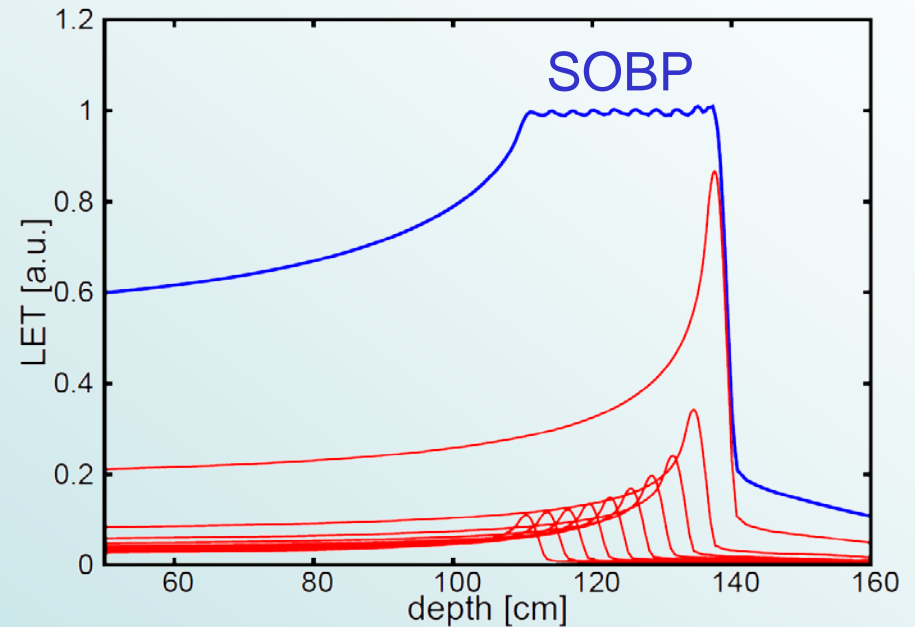
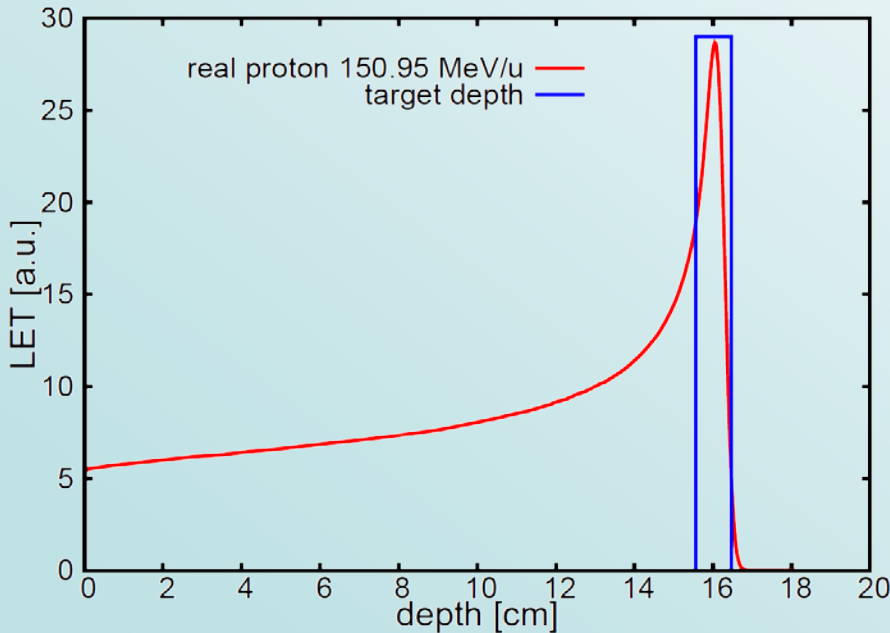




# Raster scanning technique

(Longitudinal scan direction)

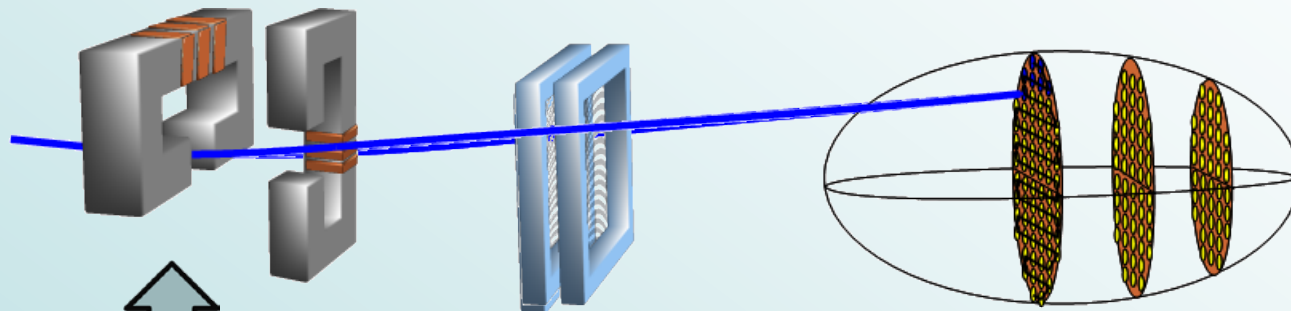
Due to the shape of the *bragg peak* not only the region where the particle stops is irradiated, but also the region before the peak (“plateau region”). This has to be taken into account during treatment planning. The dose plateau (right), result of overlapping beam with different energies, is called a *spread-out bragg peak (SOBP)*.



# Raster scanning technique

(Transversal scan direction)

In a fully active system scanning in x-y-direction (transversal to the beam) is achieved by deflecting the ion beam in a scanning magnet:



2 multiwire proportional chambers (MWPCs)

position correction

position measurement

control system

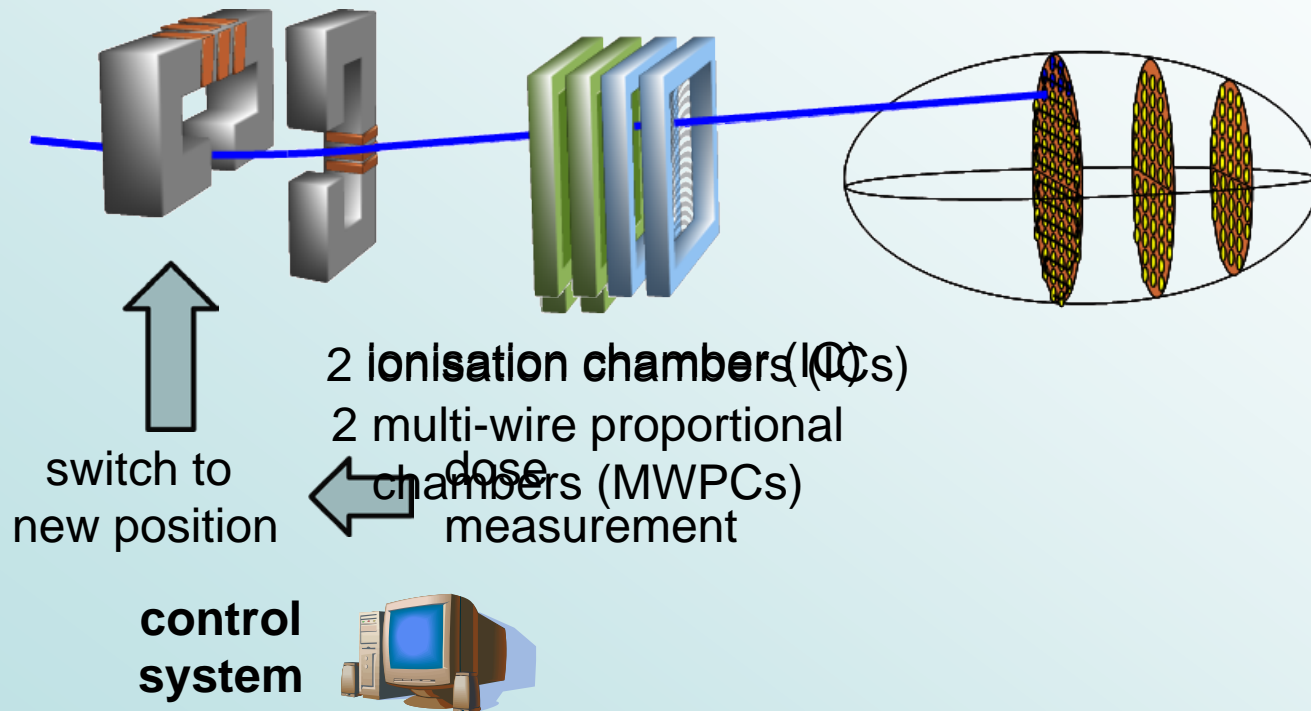


Typically, the beam will be kept at the designated position for each raster point by using a **feedback loop**.

# Raster scanning technique

(Transversal scan direction)

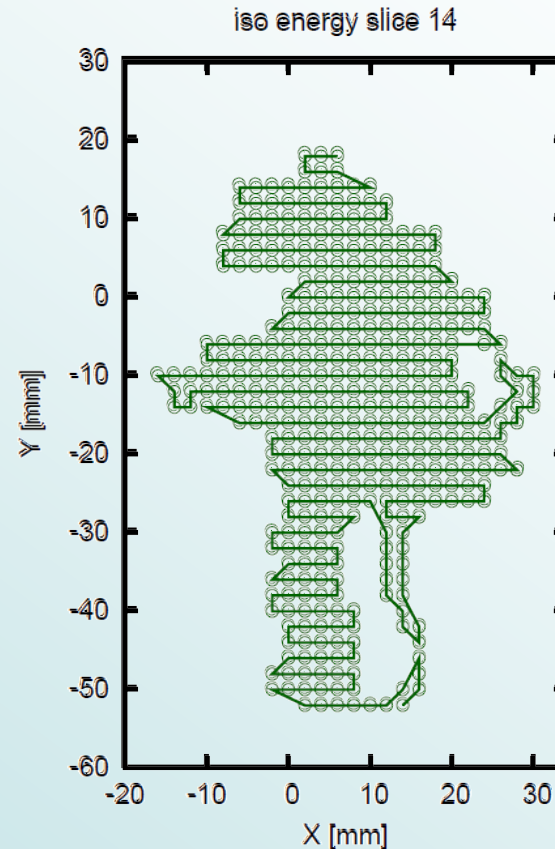
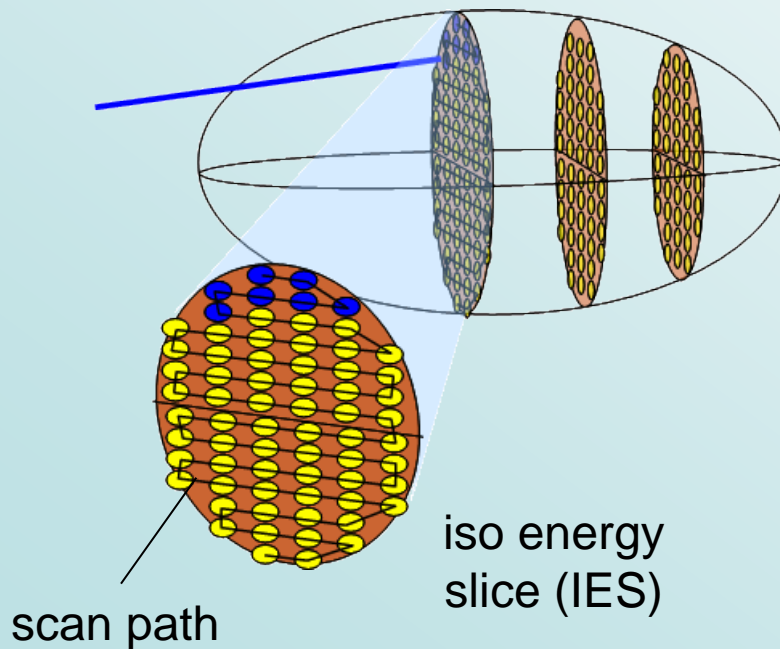
Dose controlled irradiation: the beam stays on a particular raster point until the desired dose for the point has been reached.



# Raster scanning technique

(Transversal scan direction)

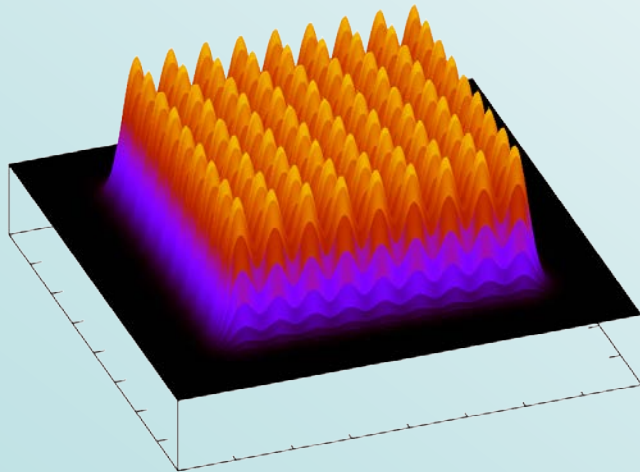
In order to cover the whole cross-section of the tumor, a **scan path** (or “work list” of points) must be defined which determines the order in which different scan spots are irradiated:



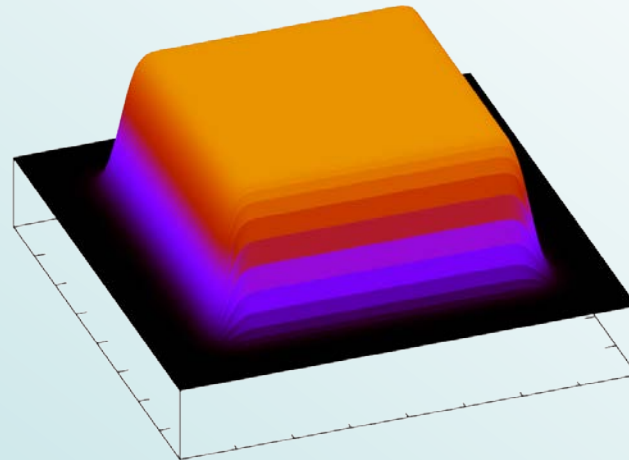
# Raster scanning technique

(Transversal scan direction)

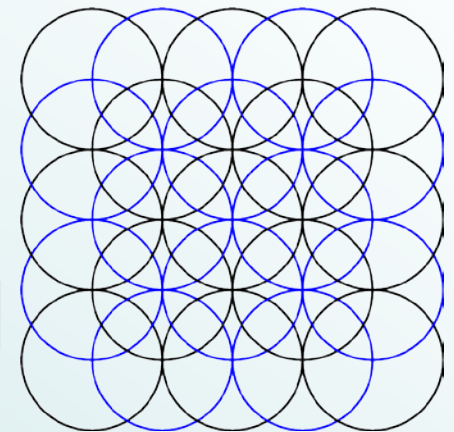
The spacing of the raster points has an influence on the dose homogeneity:



raster spacing = 3 sigma



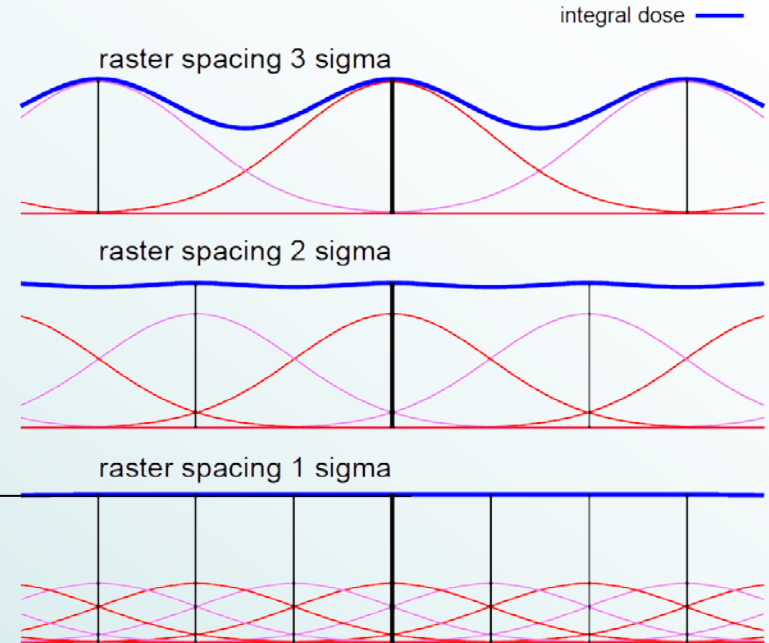
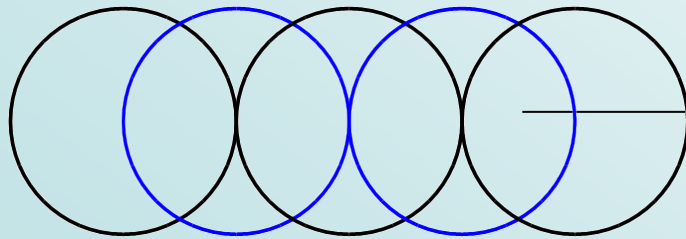
raster spacing = 1 sigma



# Raster scanning technique

(Transversal scan direction)

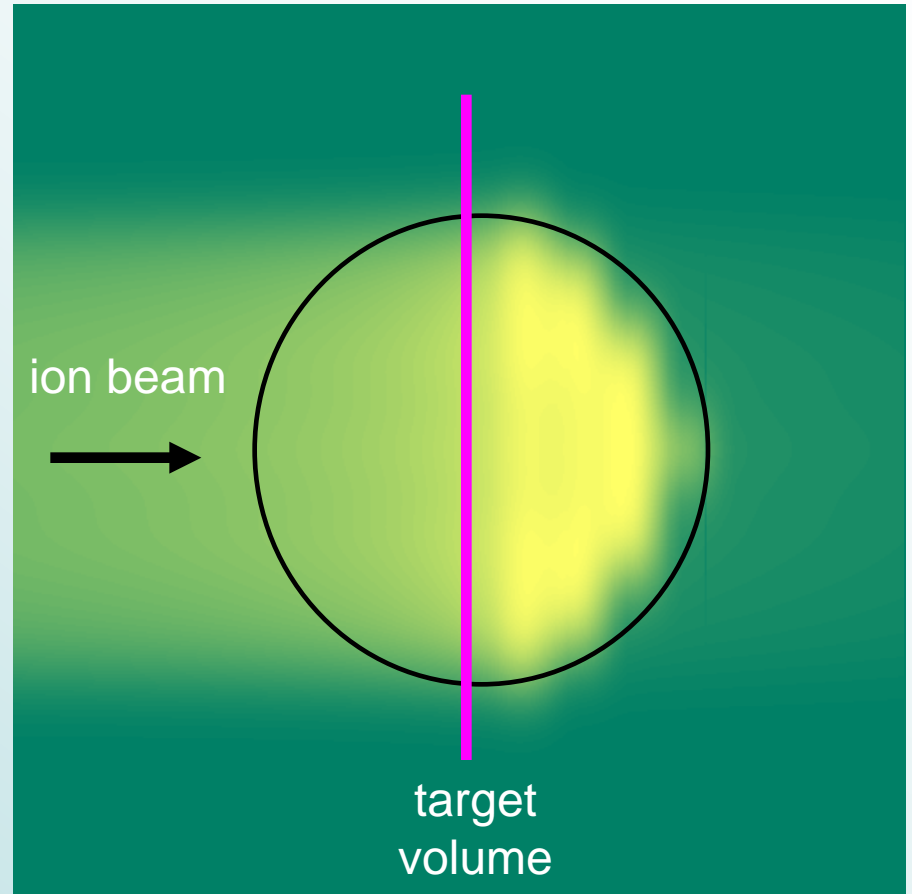
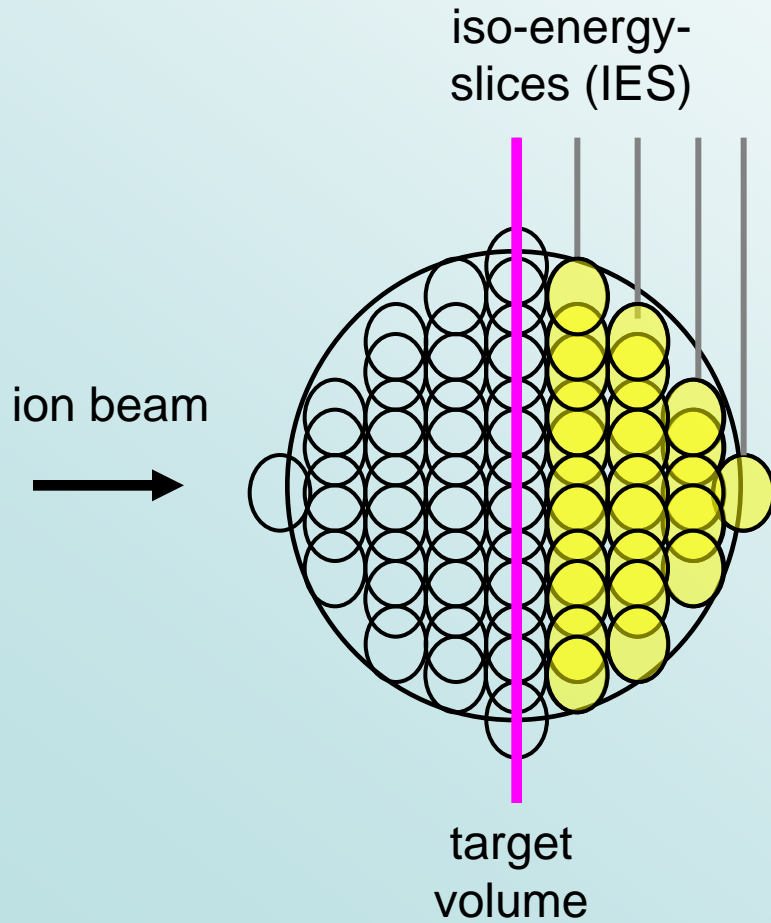
For a gaussian beam shape a dose profile will become perfectly flat if the spot separation is smaller than **one sigma**:



Dose profiles become clinically acceptable at a spacing of  $\sim 2\sigma$ ;  
For treatment planning at HIT a spacing of  $0.8\sigma$  or less is used  
as a rule of thumb to allow for some error in spot size.

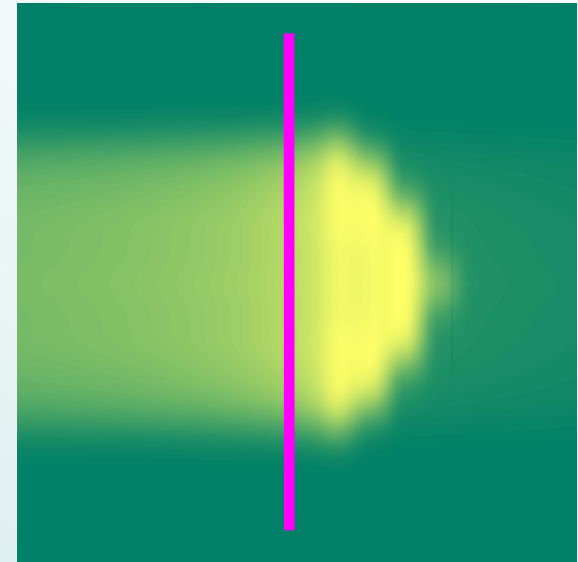
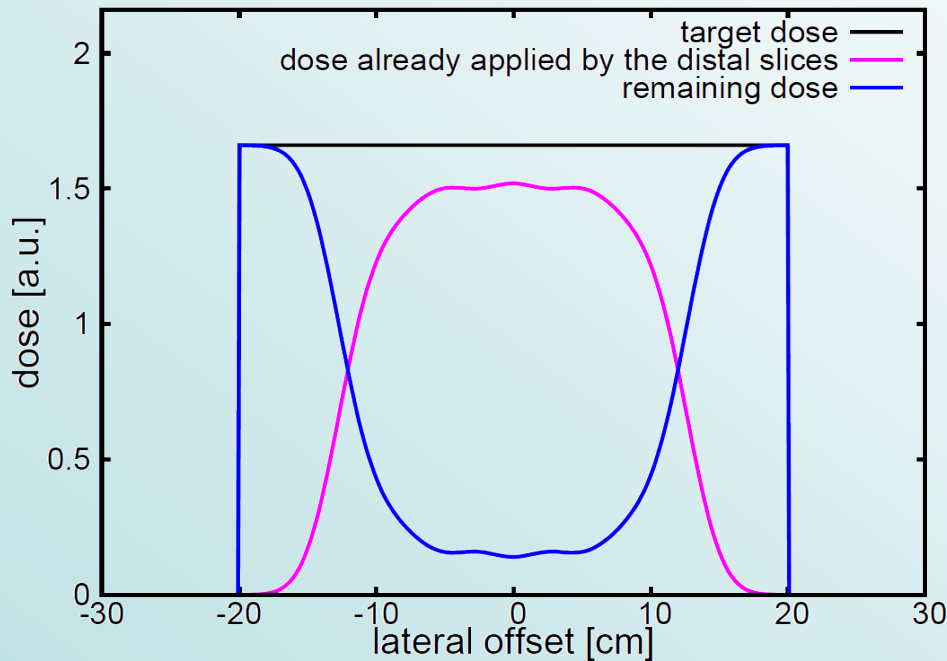
# Raster scanning technique

(Spot doses)



# Raster scanning technique

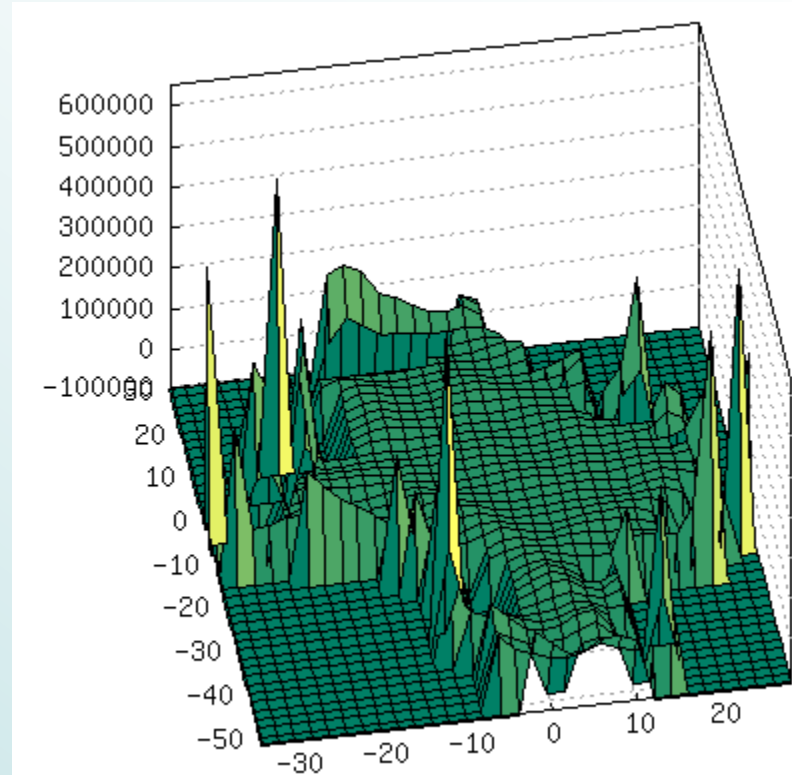
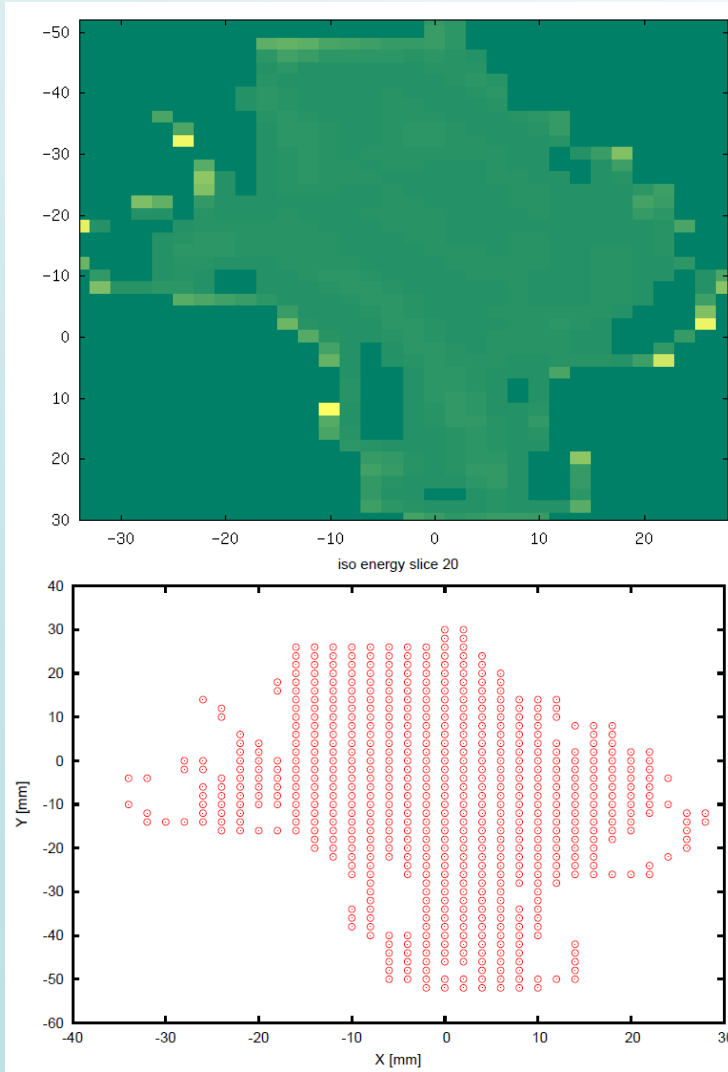
(Spot doses)



In the central region most of the dose was already applied by the distal IESs; at the lateral edges nearly the full dose is still missing.



# Raster scanning technique (Spot doses)



**The variation in particle number in one single IES can be quite large (factor of ~100)!**

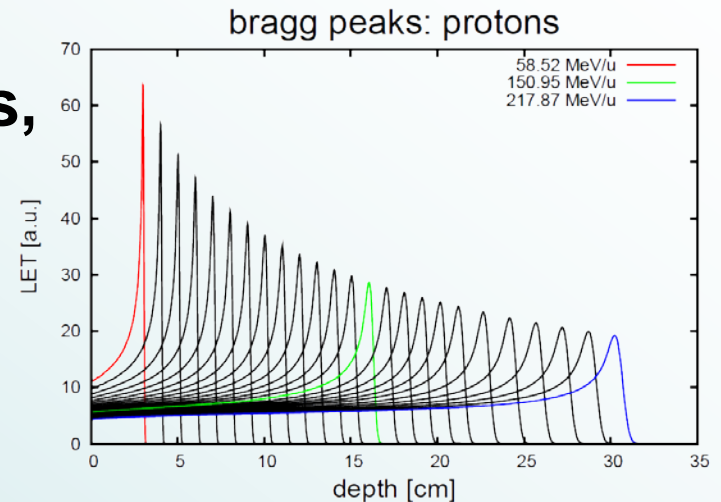
# **Interaction of Accelerator and Irradiation Technology in Particle Therapy – Examples and Challenges in Near Future**

# Raster scanning technique

(Example: HIT)

For scanning the accelerator should

- allow to request **different energies, beam spot sizes (“focus levels”) and intensities**
- deliver a different combination of all of these beam parameters **within a few seconds**
- deliver all of these combinations with a **high beam quality** sufficient for medical use
- provide a **spill pause functionality** (in case of disjointed raster point “isles” in one IES)



# Raster scanning technique

(Example: HIT /Parameters)

## Beam parameters of the Heidelberg Ion Beam Therapy (HIT)

Parameter	
ions	protons and carbon, (oxygen) – <b>2 ion sources</b> In addition: helium (3. ion source; permission for patient treatment by end of 2019 awaited)
intensity	$2 \times 10^6/s$ to $8 \times 10^7/s$ for carbon $8 \times 10^7/s$ to $4 \times 10^8/s$ for protons <b>10 steps</b> ; maximum extraction time 5 s
energy	88-430 MeV/u for carbon 50-221 MeV/u for protons <b>255 steps</b> , 1-1.5 mm spacing, 2-30 cm range in water
focus	3.5-13 mm FWHM for carbon 11-33 mm FWHM for protons <b>4 steps</b>

→ a total of  $4 \times 10 \times 255 \times 4 = \mathbf{40.800}$  settings (max.) per treatment room!

*...and much more for the gantry!*

# Raster scanning technique

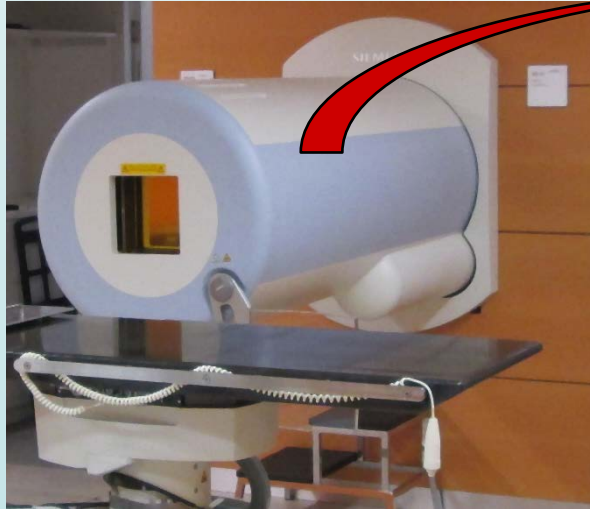
(Example: HIT / Parameters)

The therapy control system has to monitor the system on a timescale that matches the typical irradiation time of a raster point (~1-100 ms); this implies:

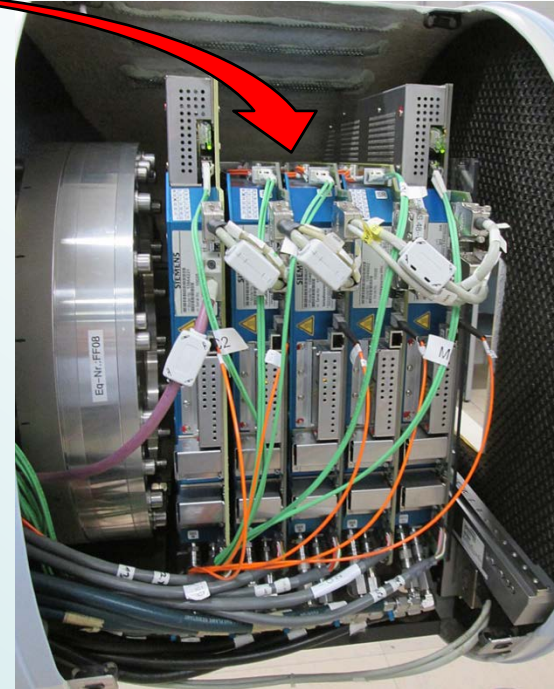
- Dose controlled irradiation system: dose measurement on a time scale that is short compared to the irradiation time of one point (here: 10  $\mu\text{s}$  – faster than drift time of ions in the ionization chamber);
- Position measurement with MWPCs on a time scale that is “short” compared to typical position variations (HIT: 250  $\mu\text{s}$ )
- Beam position feedback system (linear) – see next two slides
- Online monitoring of beam focus size and limit checks; focus feedback loop planned with next generation control system – beam optics have to be considered!

# Raster scanning technique

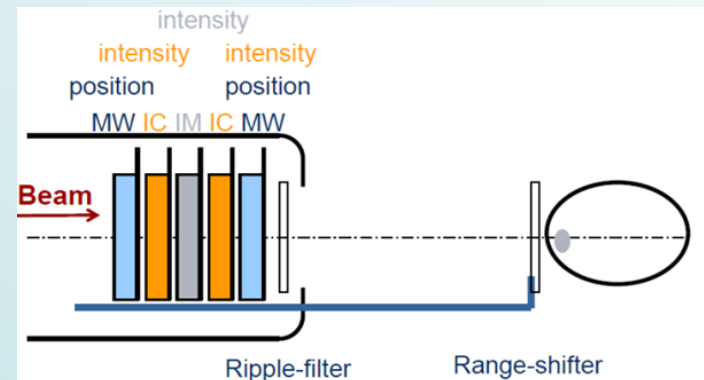
(Example: HIT / BAMS)



Beam  
Application  
and  
Monitoring  
System

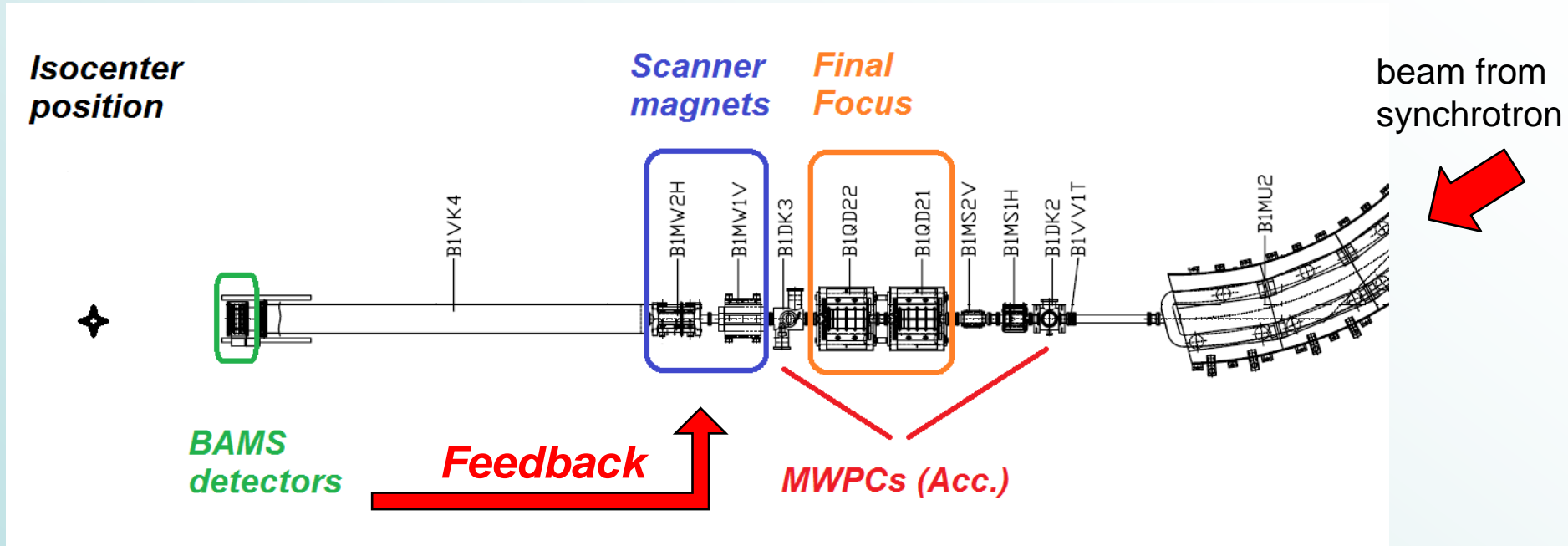


- 2 (redundant) MWPCs for **profile monitoring** and **position feedback**
- 3 (diverse redundant) ionization chambers for **dose measurement**



# Raster scanning technique

(Example: HIT / beam position feedback)



The „source point“ for the raster scanning system (example: horizontal treatment room at HIT) are the scanner magnets, where the beam gets small angles (h,v) to hit the “right” voxel position in the isocenter plane (and a corresponding position in the BAMS detectors). In case of deviations (h,v) the scanning angles are corrected, the beam optics simply follows the intercept theorem.

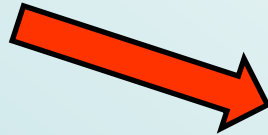
# Combining PT and MRI



# Combining PT and MRI

## Motivation:

- “Seeing what you treat” → Online diagnostics would be favorable
- CT is almost standard today, but MRI causes no further radiation dose (especially important in pediatric treatments!)
- Tumor shrinkage during therapy – avoidance of errors in adapted dose allocation

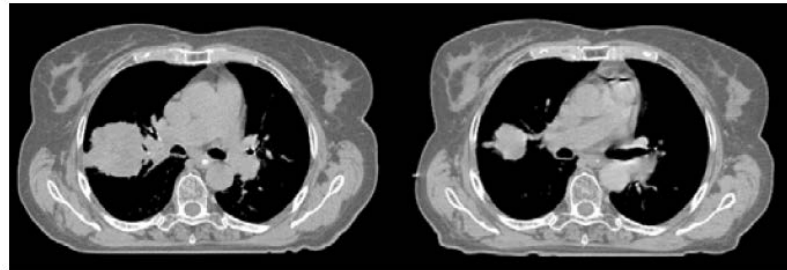


- MRI-Linac Systems (with photon beams) are currently being introduced in radiotherapy, see slide after next

## Lung Tumor shrinkage during p-therapy

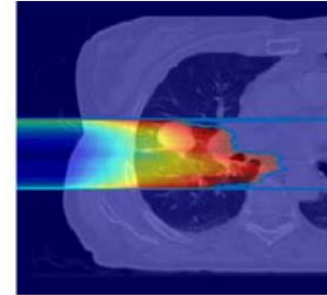
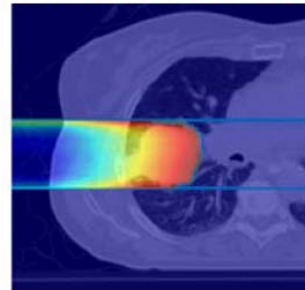
Initial Planning CT  
GTV 115 cc

5 weeks later  
GTV 39 cc



Beam stops at distal edge

Beam overshoot



Courtesy of S. Mori, G. Chen, MGH, Boston

# Combining PT and MRI

## Furthermore:

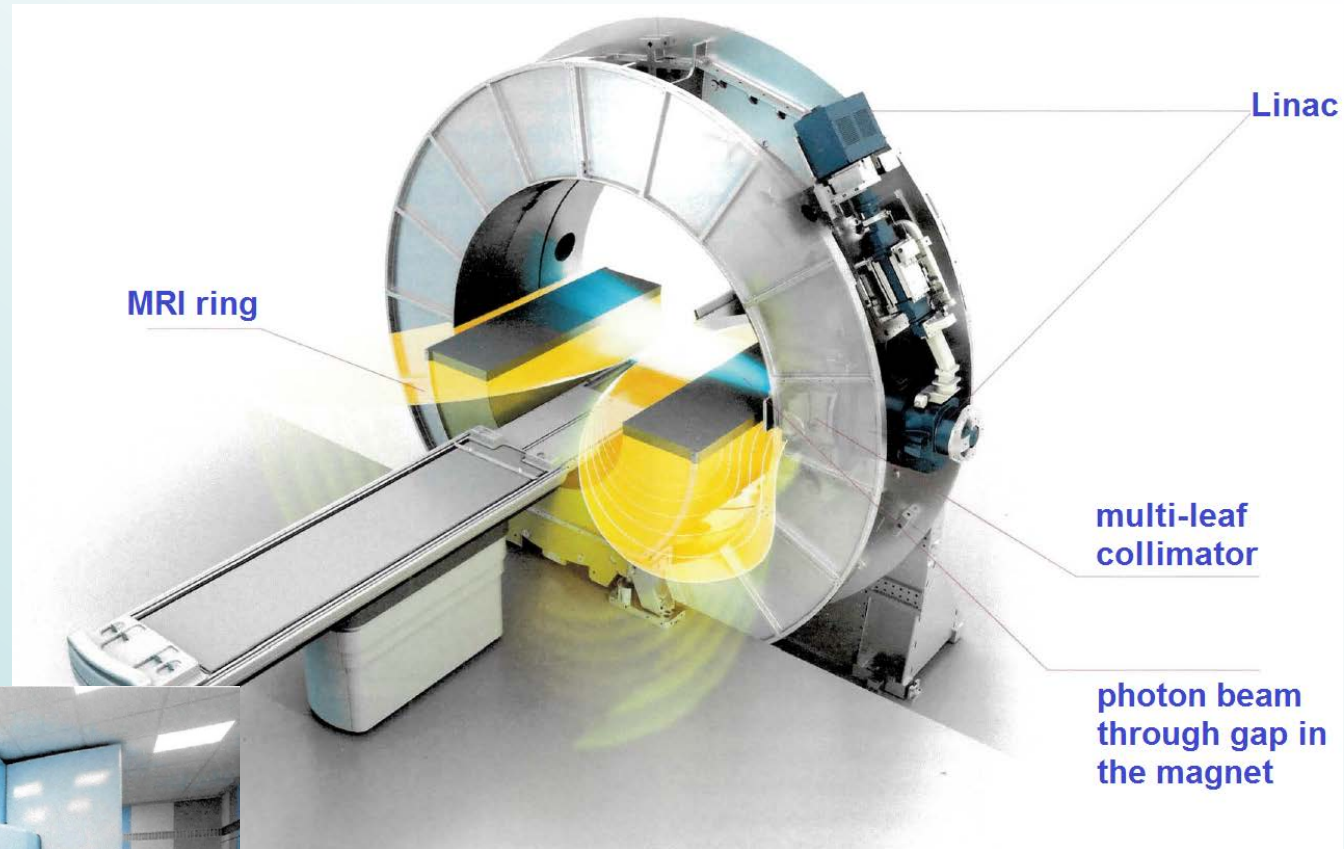
Doctors normally use large margins around the tumor volume because of possible range uncertainties, movement of the tumor along the irradiation, etc. → Better diagnostics is necessary to shrink these margins!



A 6.5 mm thick margin (peel) consists of the same volume as a 5 cm diameter target (orange).

# Combining PT and MRI

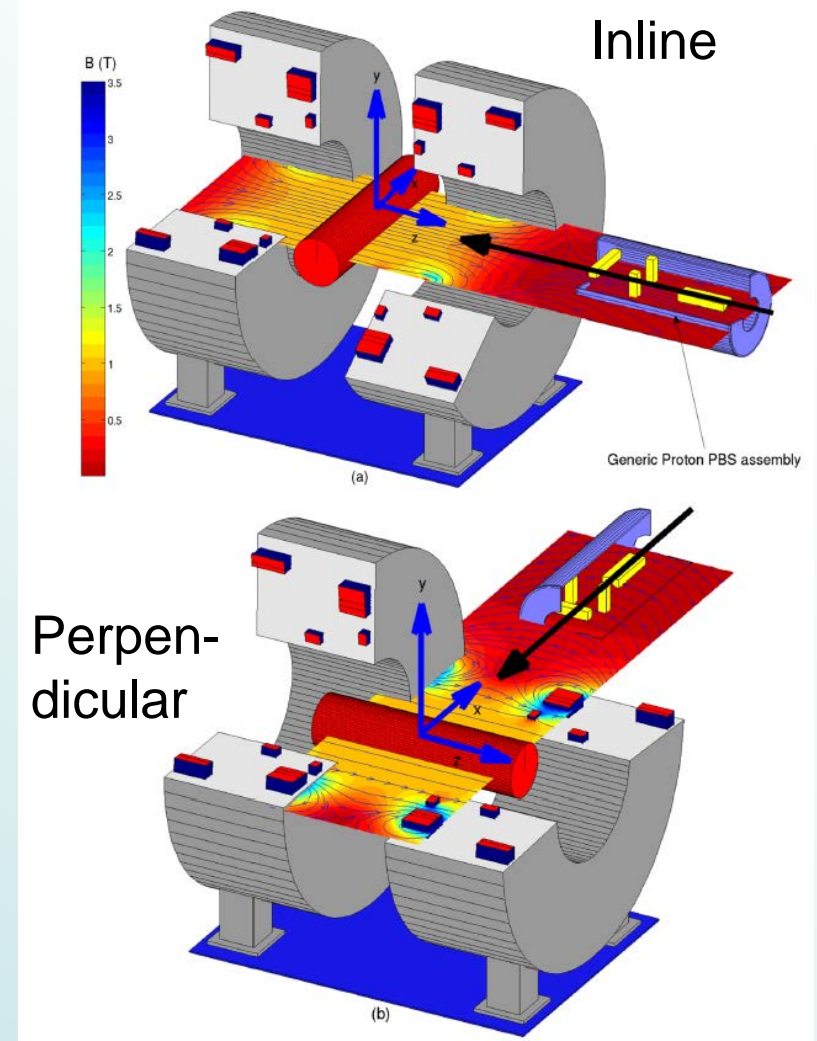
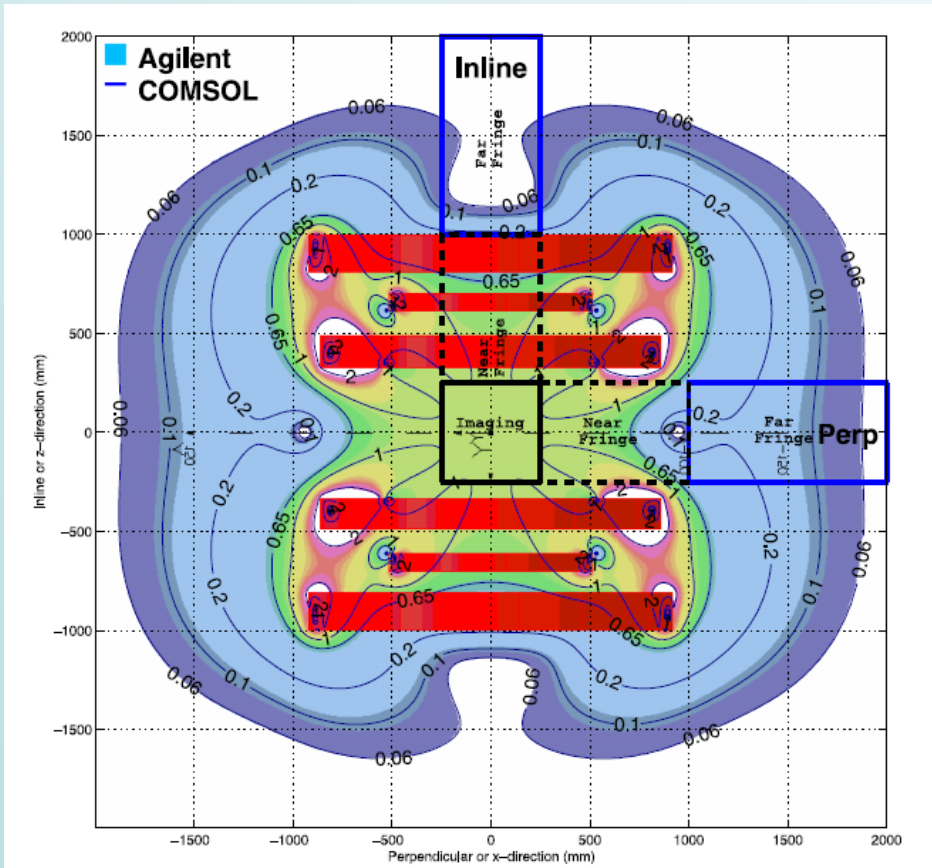
Cross section  
of a combined  
MRI and  
photon linac



Installation of the MRIdian® Linac  
(with 0.35 T magnetic field) at the  
University Hospital Heidelberg /  
Section RadioOncology

# Combining PT and MRI

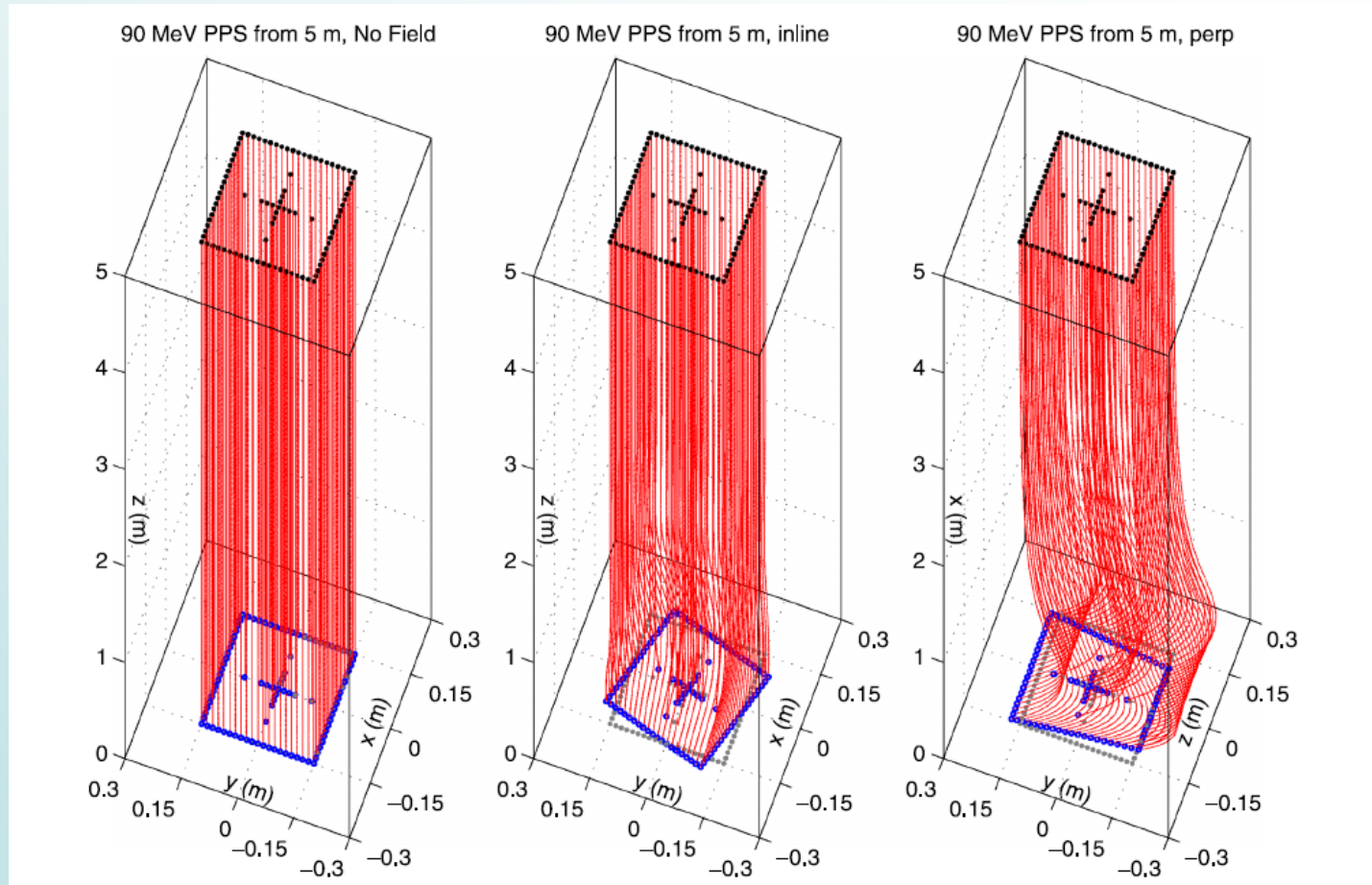
MRI devices have complex magnetic fields, which would interact with proton/ion beams – compensation, feedback?



Possible arrangements of MRI device vs. beam direction

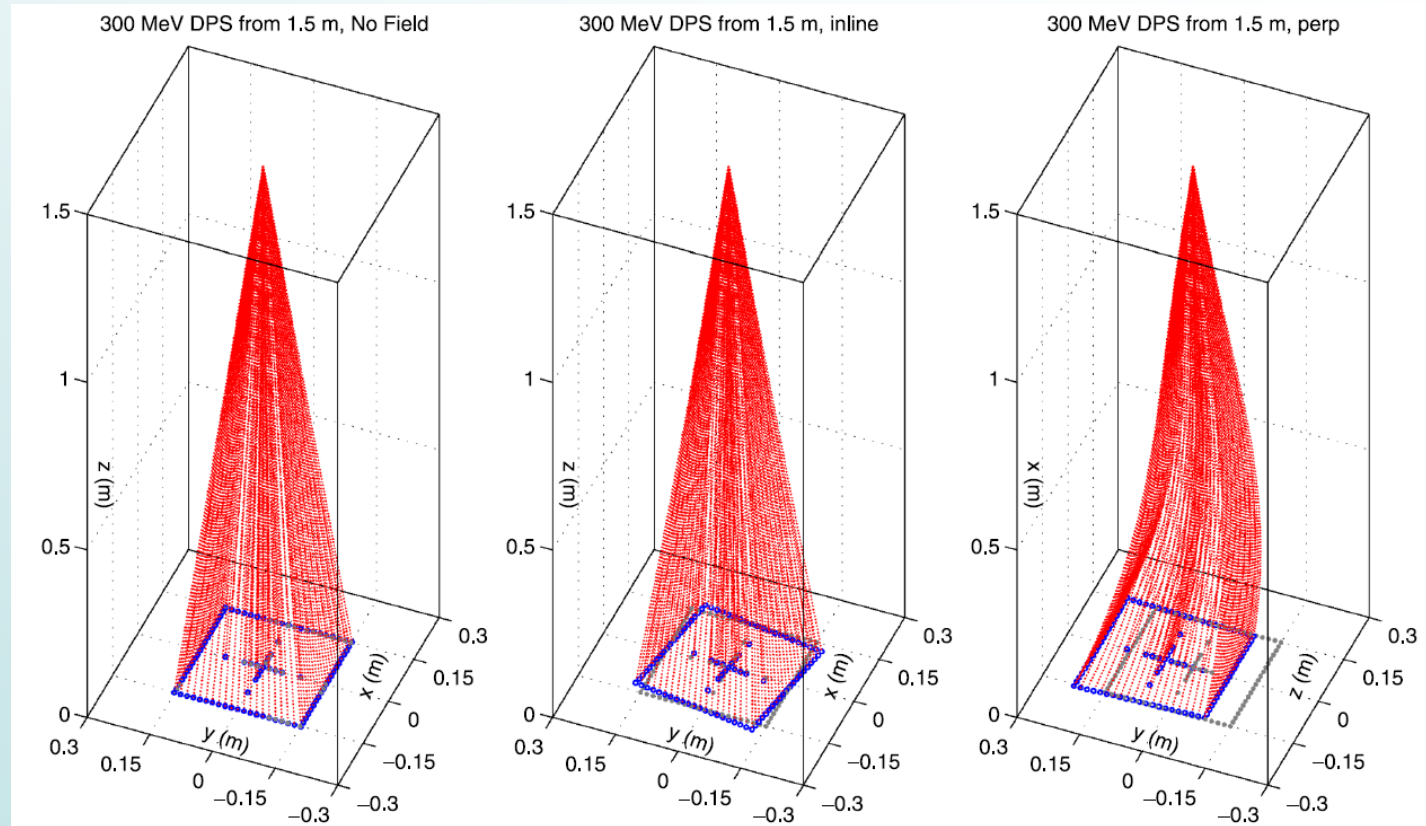
# Combining PT and MRI

In case of a parallel particle source (PPS) the MRI device causes (complex) distortions and deflections of the beam!



# Combining PT and MRI

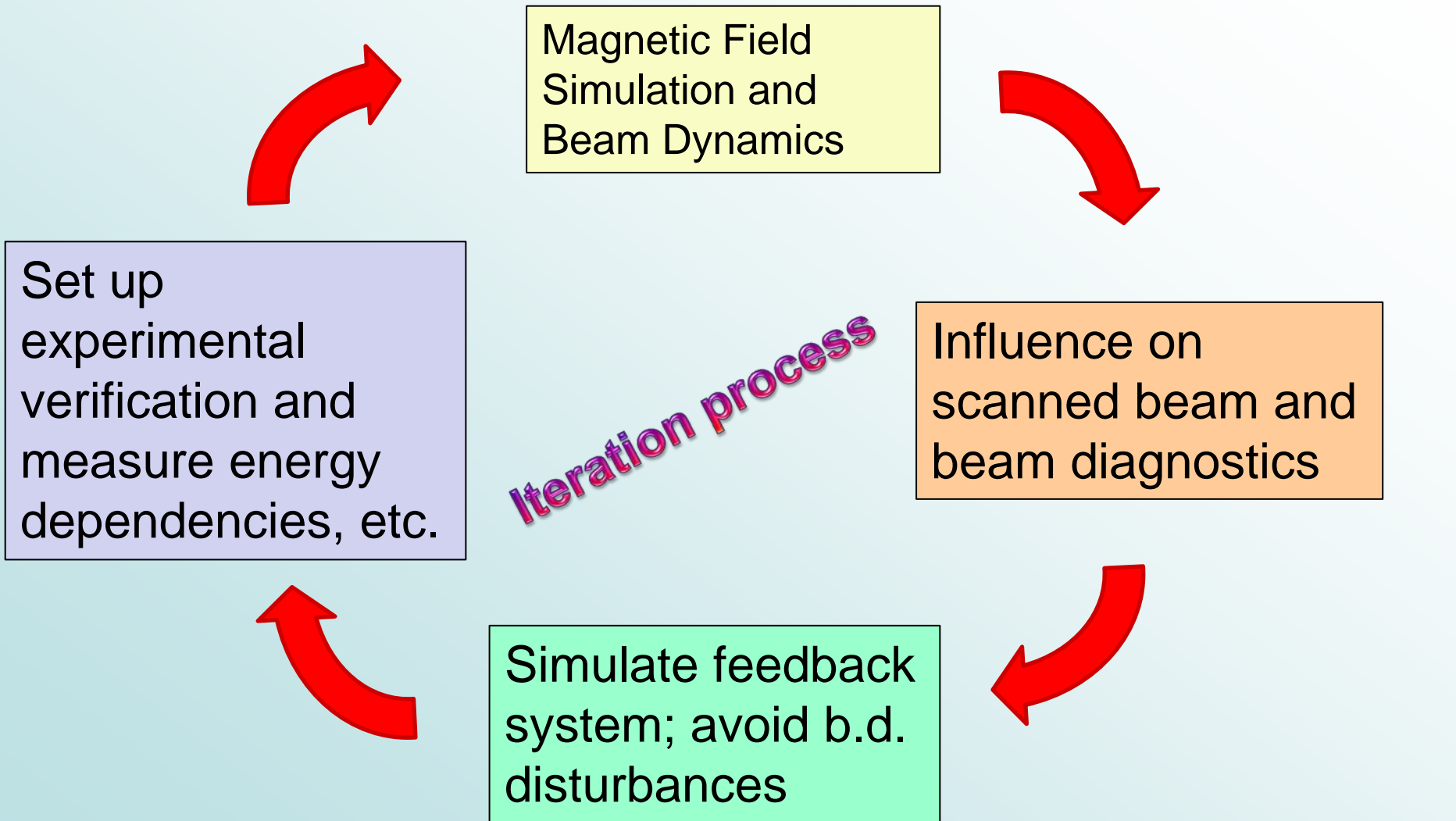
In case of a Diverging point source (DPS), e.g. a raster scanning system, similar (complex) distortions and deflections of the beam occur!



## Furthermore:

- Effects on secondary particles have to be taken into account!
- Influence of pulsed magnetic fields (MRI) have to be carefully examined!

# Combining PT and MRI



***Solutions still under study...***

# Dynamic Intensity Control – a Variable Feedback System for Fast Dose Application



# Dynamic Intensity Control

Motivation:

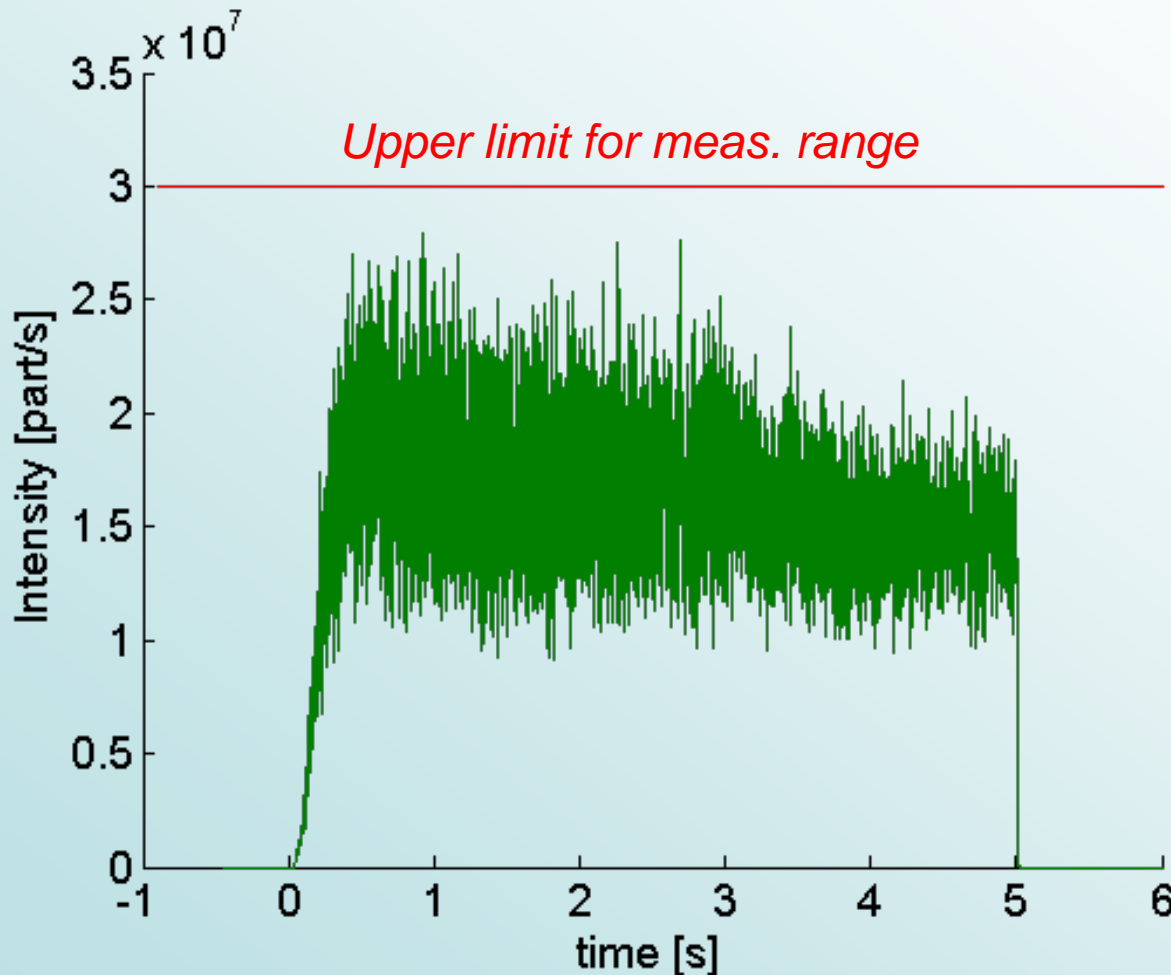
Therapy works fine – why do we need higher performance in dose application?

Reasons for the reduction of individual treatment time are:

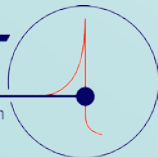
- Higher patient comfort (locally immobilized!)
- Higher dose conformity
- More patients
- Economic facility operation



# Dynamic Intensity Control

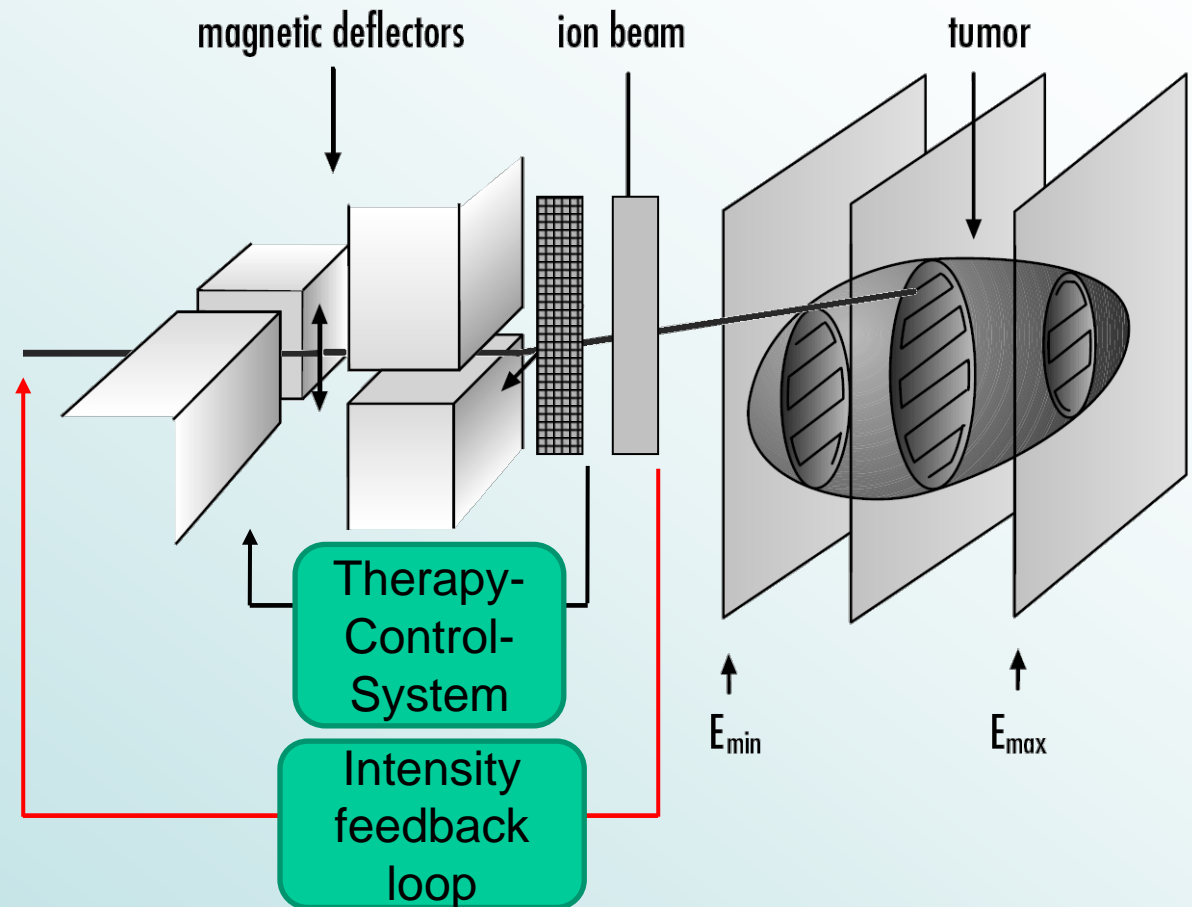


- Typical spill structure achieved with RF knock-out method
- Ideally intensity as high as possible close to the limit
- Reality: Scanning velocity is lower than desired
- **Spill-quality is essential for the treatment time!**



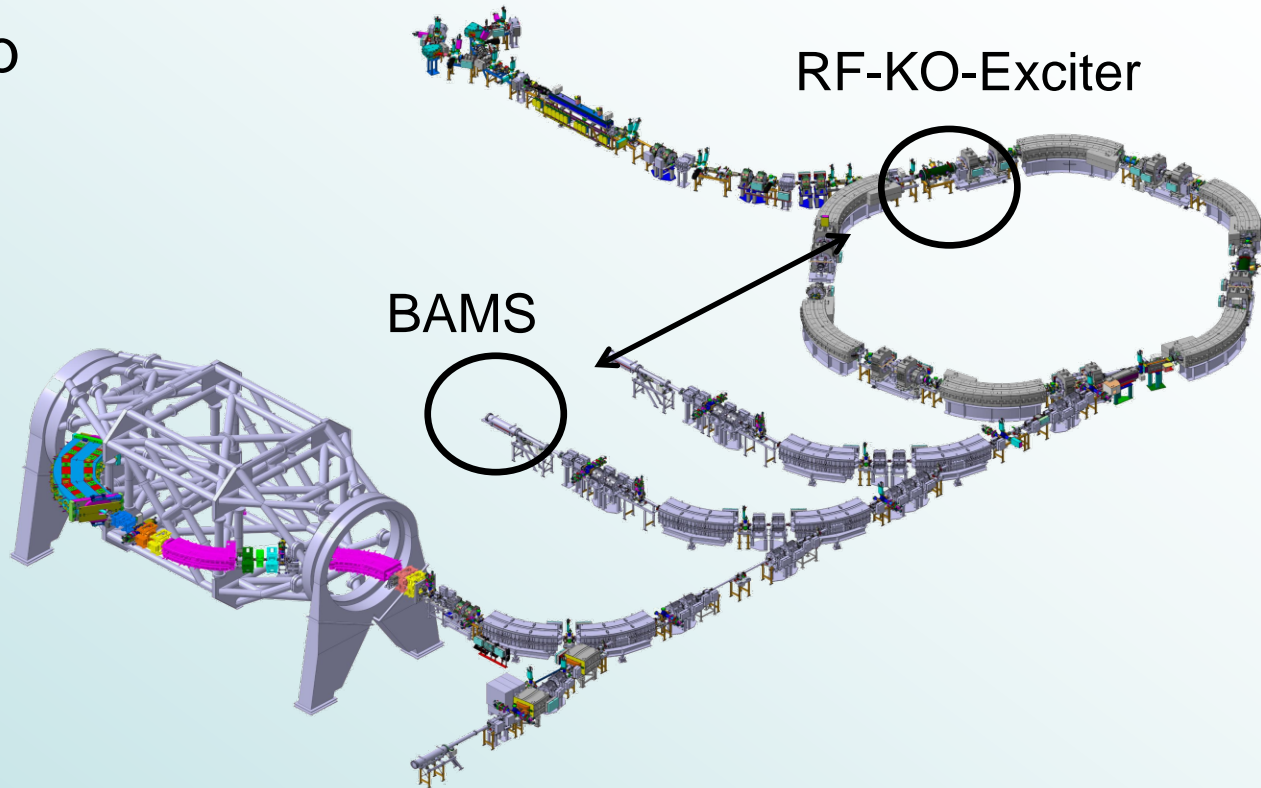
# Dynamic Intensity Control

- Use intensity signal
- Add feedback loop
- Position and intensity is measured, the scanning velocity **and the intensity** is adapted
- Challenge: Coupling the *medical product* with the *industrial product* accelerator



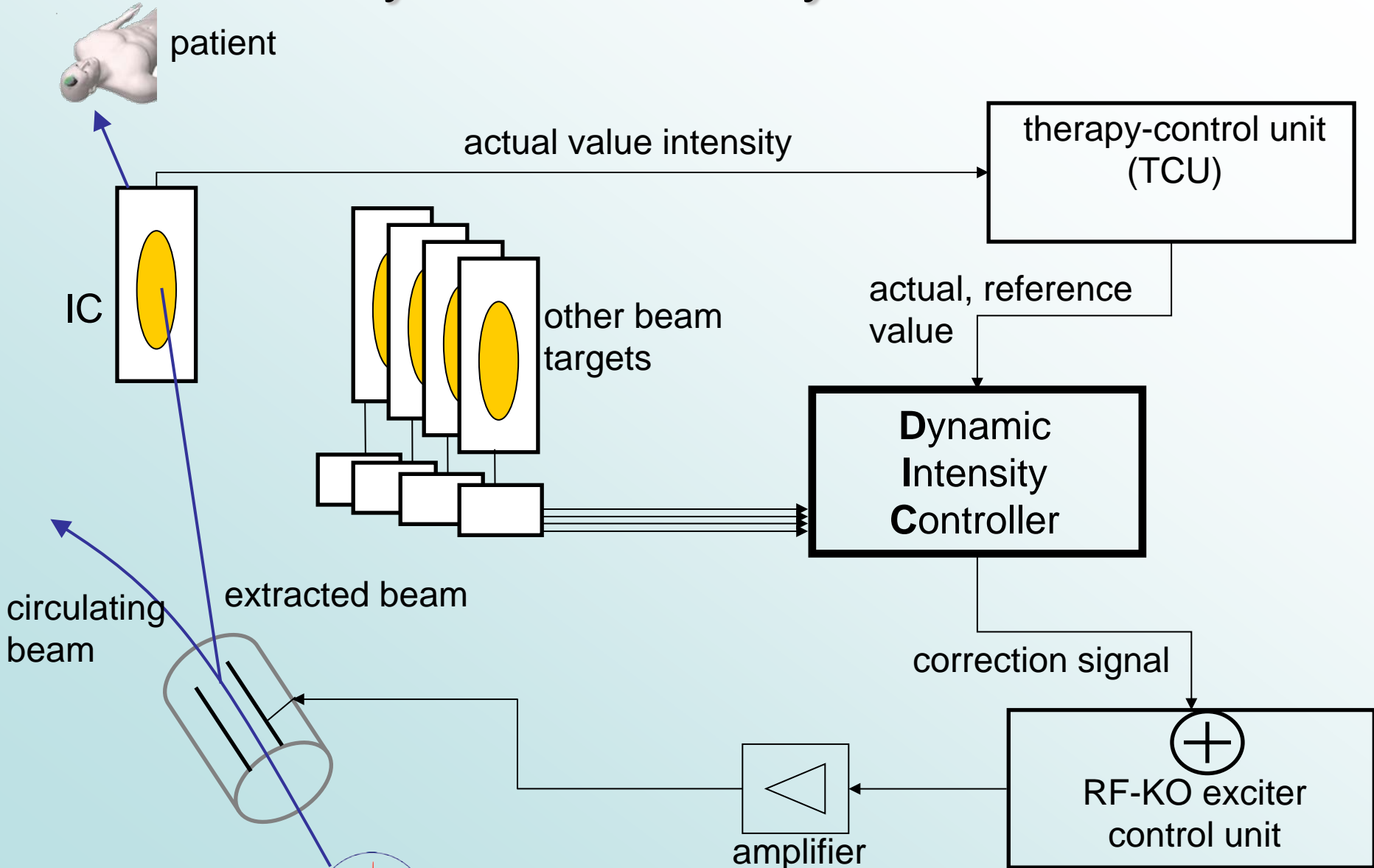
# Dynamic Intensity Control

- Use intensity signal
- Add feedback loop
- Position and intensity is measured, the scanning velocity **and the intensity** is adapted
- Challenge: Coupling the *medical product* with the *industrial product* accelerator



4 treatment rooms  
plus beamdump  
with IC detectors

# Dynamic Intensity Control



# Dynamic Intensity Control

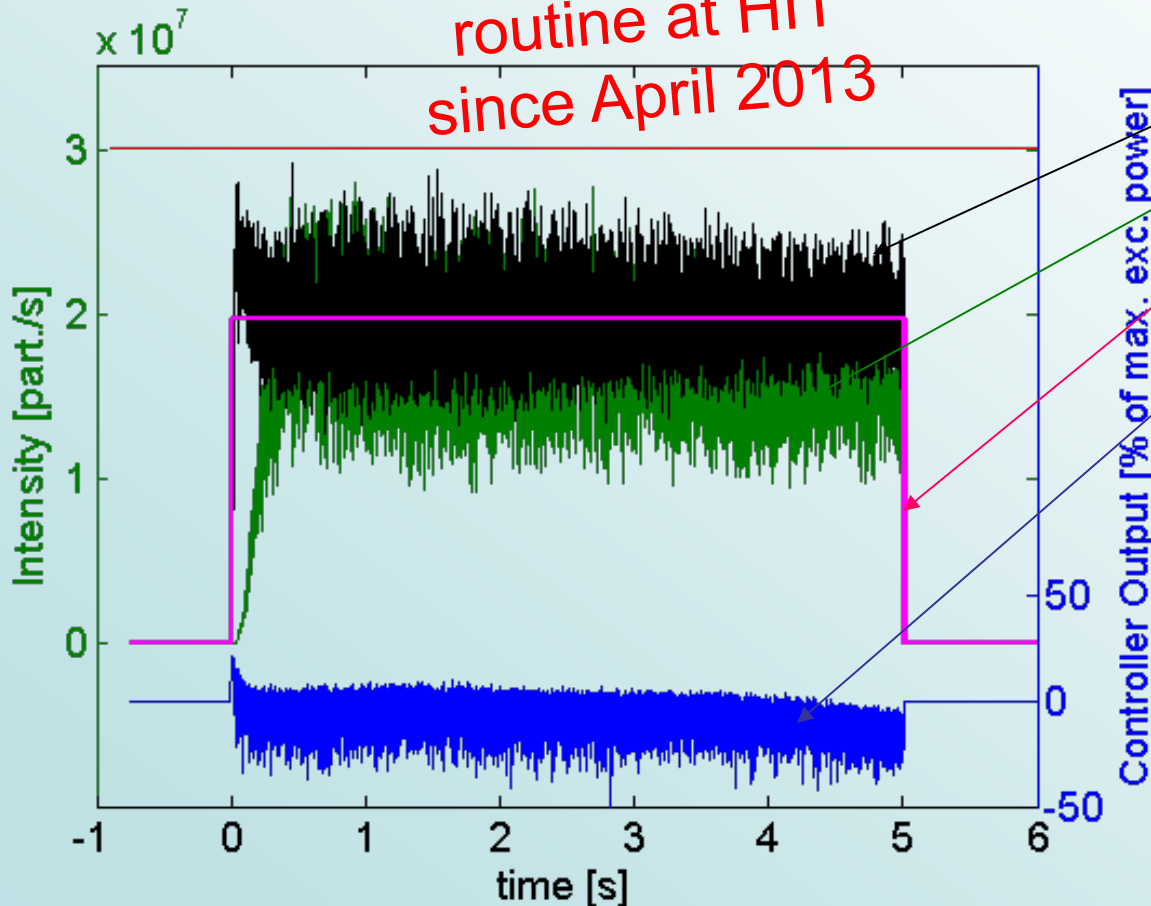
## Feedback realization:

- PID controller
  - "P" for a fast response
  - "I" for no remaining control deviation
  - "D" optional and currently deactivated
- PID-parameters as a function of (ion, energy, intensity)
  - $\approx 10000$  combinations
  - 1% was defined in commissioning, all other values were calculated by spline interpolation
- Additional features:
  - Mitigate intensity overshoot
  - "Early abort" – controller realizes when synchrotron is empty
  - ...

# Dynamic Intensity Control

## Results (Step1):

In clinical  
routine at HIT  
since April 2013

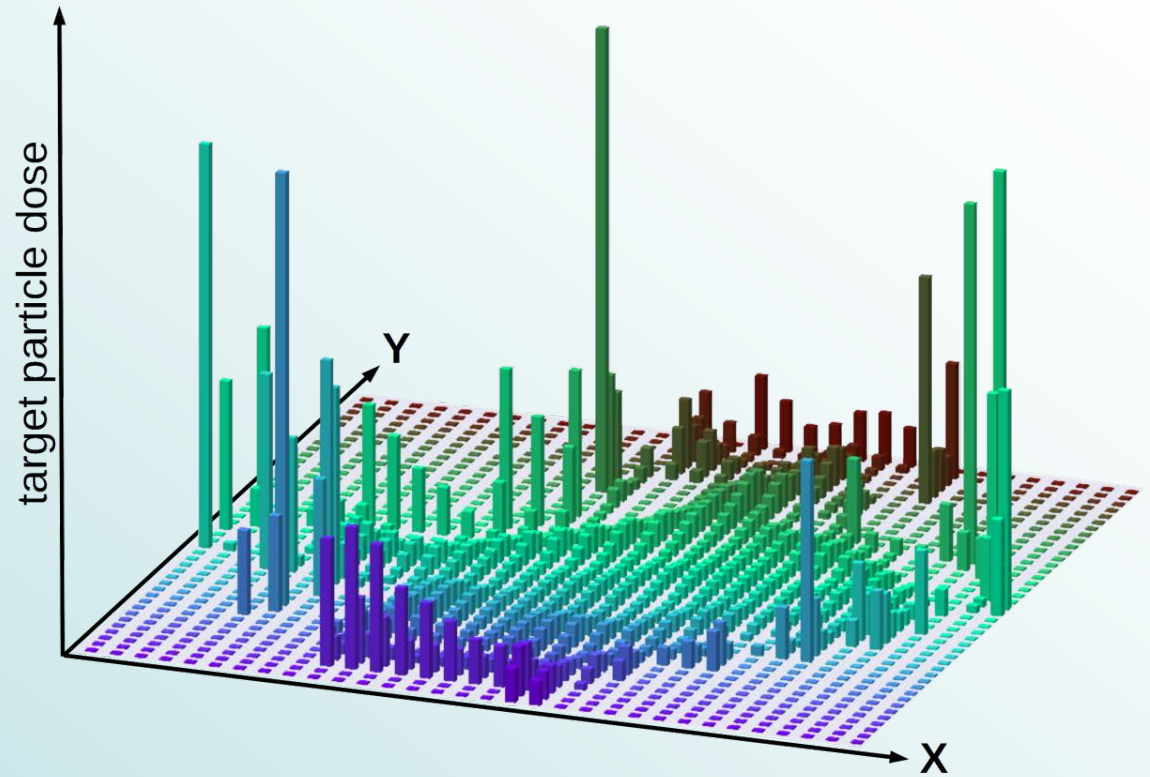


- Spill shape
  - With feedback
  - Without feedback
  - Reference value
  - Controller output

- Higher average intensity (15 %!)
- Faster irradiation
- Less machine tuning

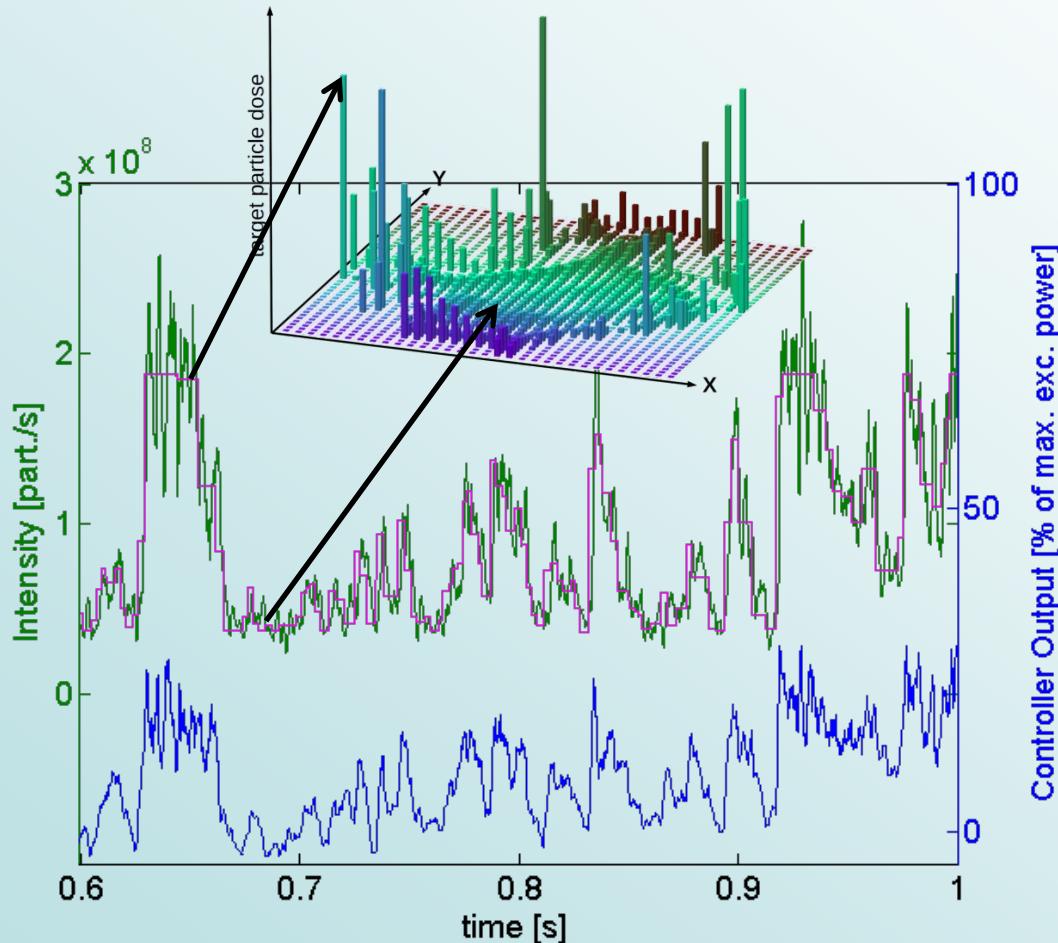
# Dynamic Intensity Control

- Example of dose distribution of one slice
- Lowest particle fluence determines intensity for whole slice
- Fixed intensity: irradiation time per raster point can vary by a factor of  $> 100!$





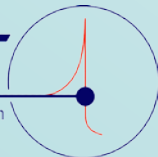
# Dynamic Intensity Control



2<sup>nd</sup> Step of DIC development:

Intensity-modulated spill

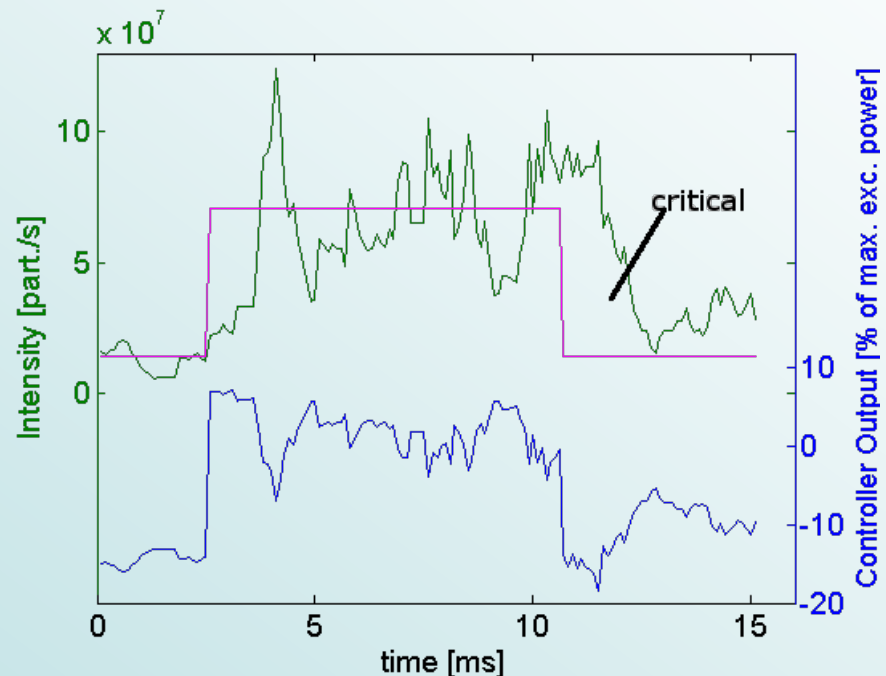
- Rasterpoint individual **reference value**
- Feedback loop adapts the **actual intensity**
- Beam-on time can again be reduced by **≈45%!**
- **In clinical operation at HIT since April 2014!**



# Dynamic Intensity Control

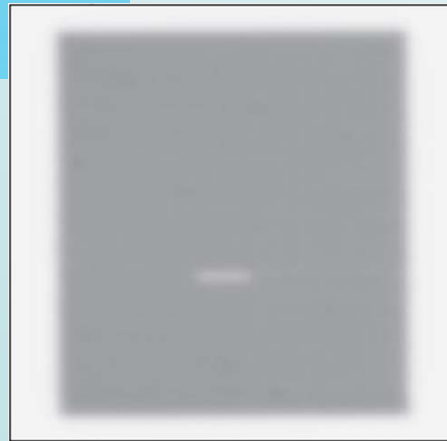
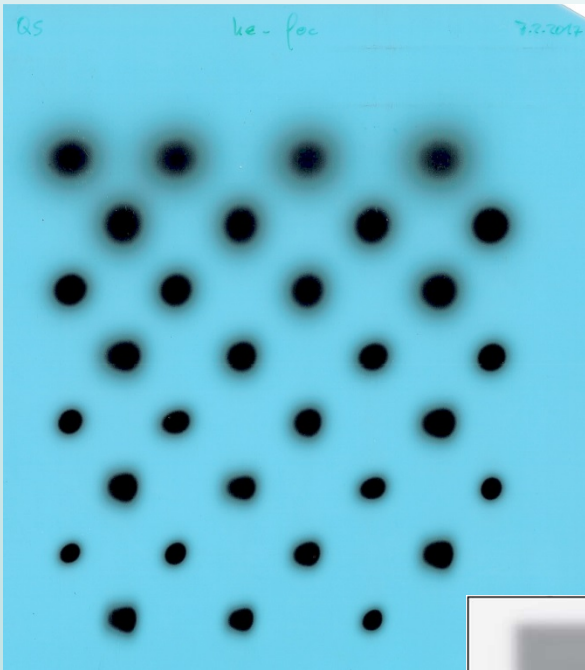
## Challenges and limitations:

- Limits of the feedback loop due to dead times
  - Signal detection and digitization, ionization chamber  $\sim 150\mu\text{s}$
  - Particle excitation  $\sim 0 - 600\mu\text{s}$
  - Latencies in digital transmission  $\sim 100\mu\text{s}$
- Irradiating too fast leads to interlocks and must be avoided!
- Reference value pattern must be defined in an intelligent way!



# Using Scintillating Screens for Beam Quality Assurance

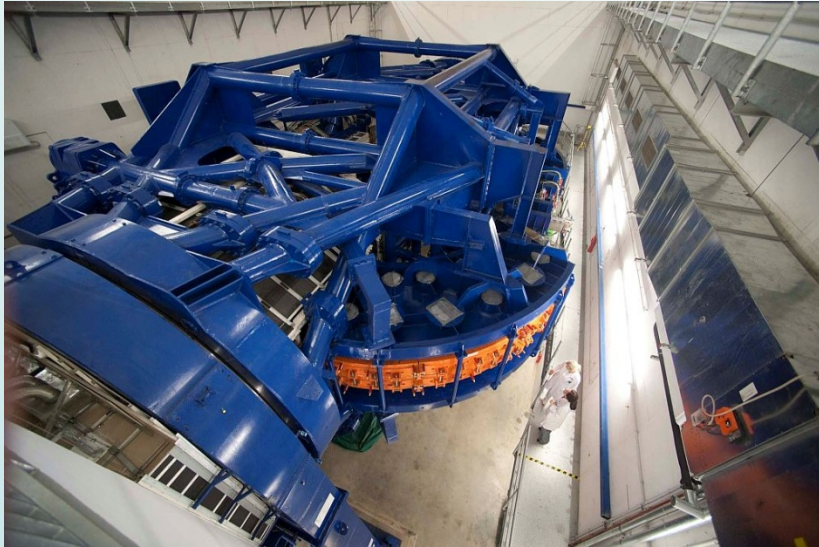
# Beam Diagnostics for QA



Not long ago irradiated films were produced and evaluated for the daily QA by the medical physicist to check:

- Focus width
- 2D „Roundness“ of the beam
- Homogeneity of dose application

And in the case of a gantry this for a lot of angles of incidence...

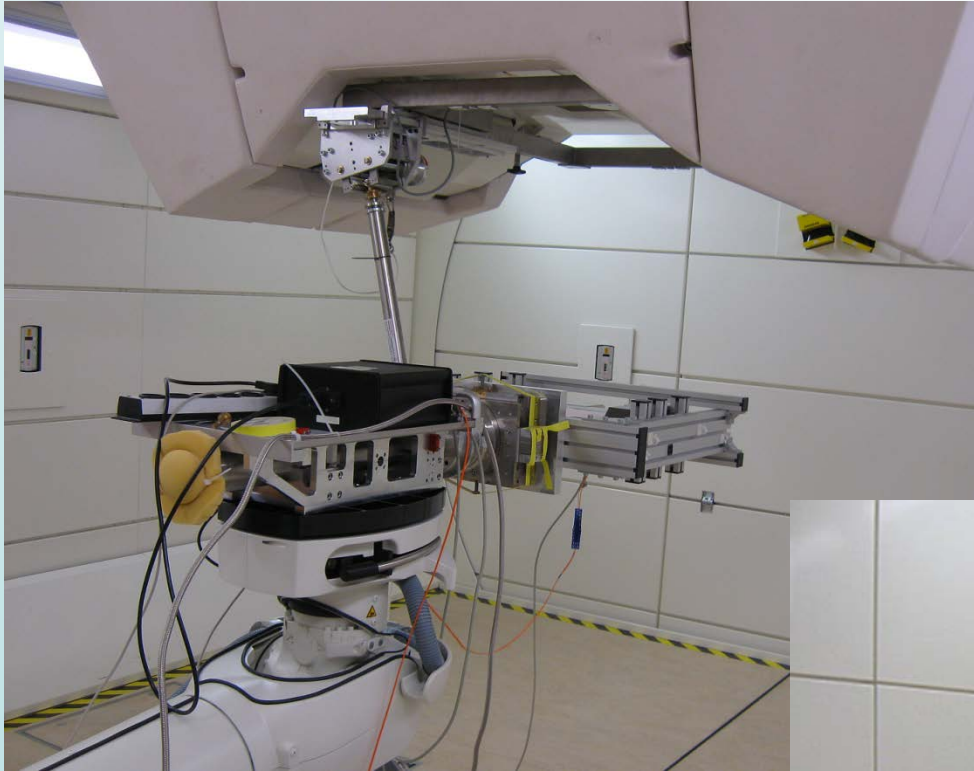


## The HIT gantry:

- Structure with beamline (upper left)
- Treatment room (lower left)
- ...and rotating (below)



# Beam Diagnostics for QA

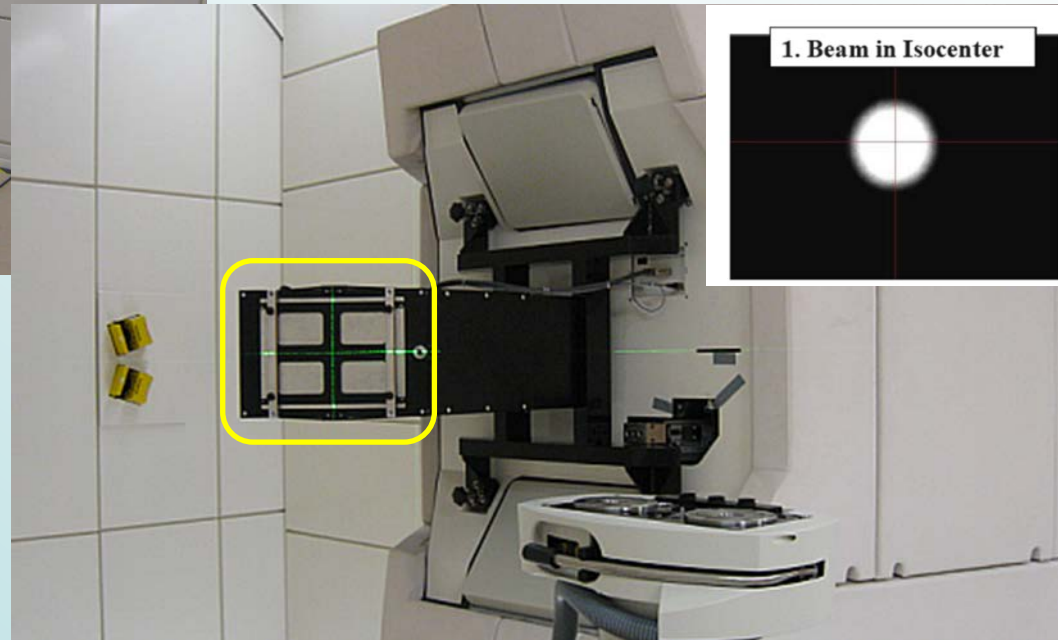


Left: Adapter to robot with rotating mount for

- Films
- MWPCs
- Other QA measurement equipment

→ **Patient coordinate system**

Right: Beam Diagnostics for commissioning – large scintillating screen fixed to gantry nozzle → **Gantry coordinate system**

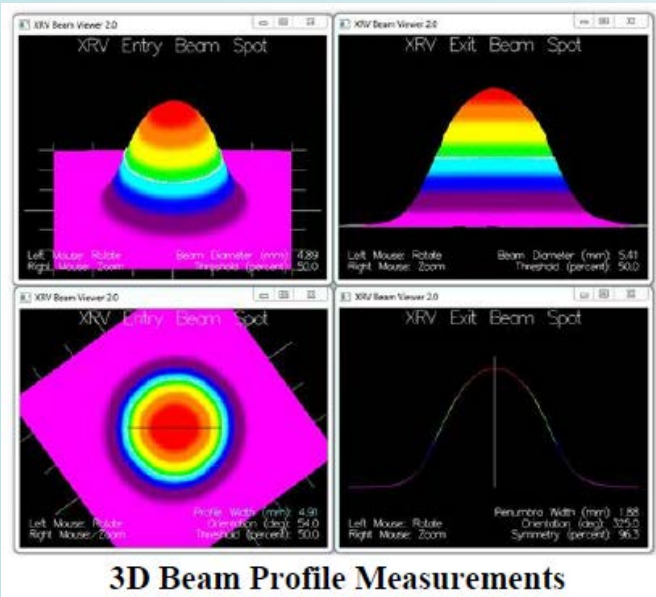
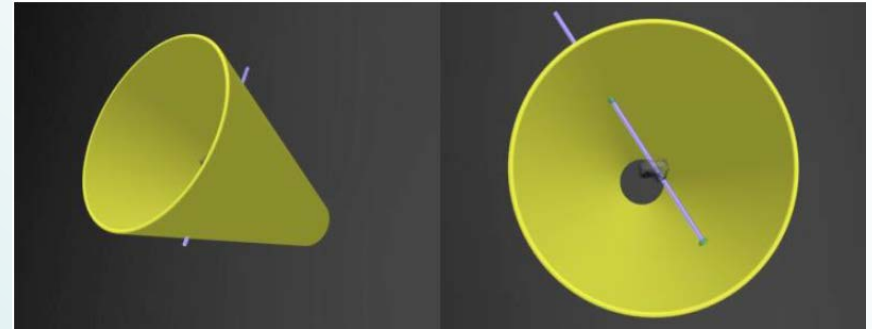


# Scintillating Screens for QA

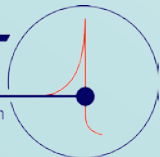
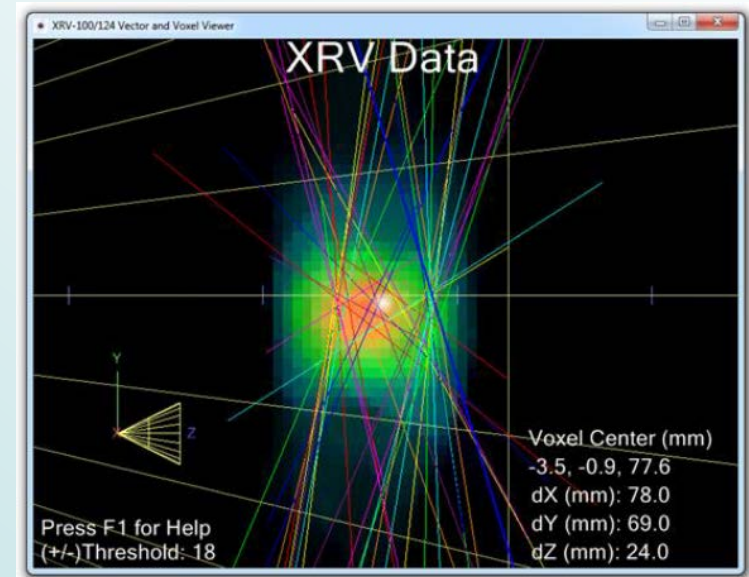
## XRV-124



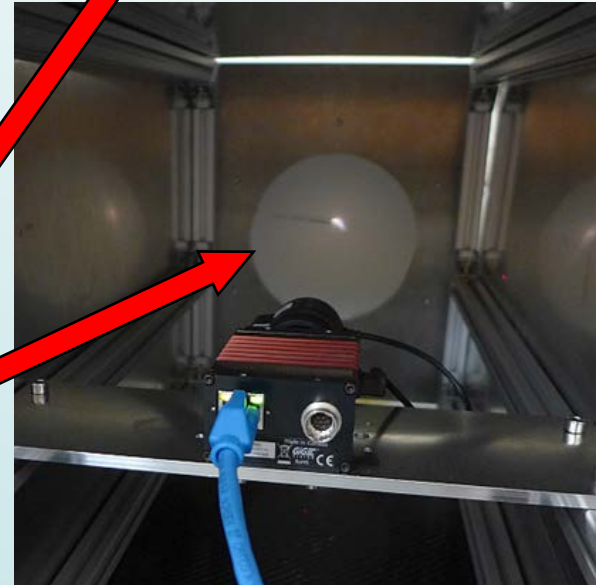
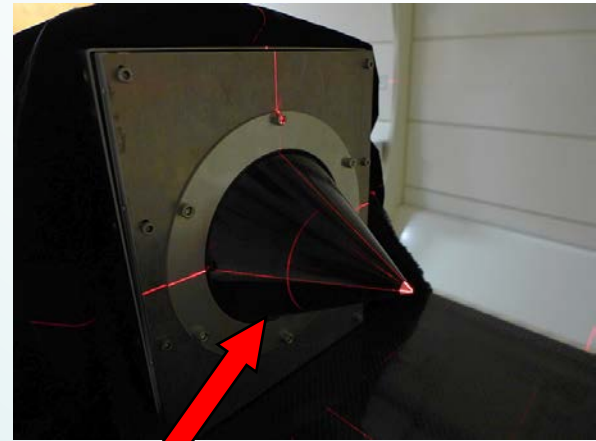
System by [www.logosvisionsystem.com](http://www.logosvisionsystem.com)  
for photon and proton beams



Geometrical reconstruction leads to beam profile data including focal point and focus width



# Scintillating Screens for QA



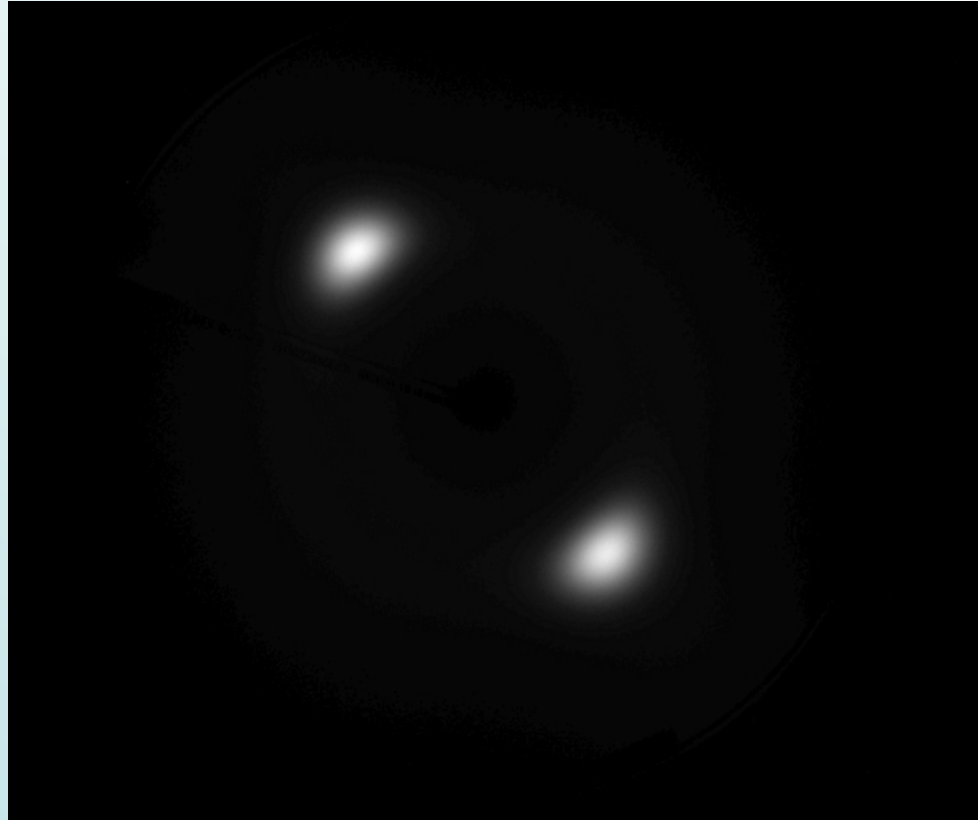
Prototype at HIT (with higher dynamics needed for ion beams); cone-shaped shell made of carbon fiber; Scintillating materials: Lanex or P43 foils from Proxivision ([www.proxivision.de](http://www.proxivision.de))



# Scintillating Screens for QA

Series of images taken shot by shot at different gantry angles:

Proton beam with 106.8 MeV and a focus width of 15 mm



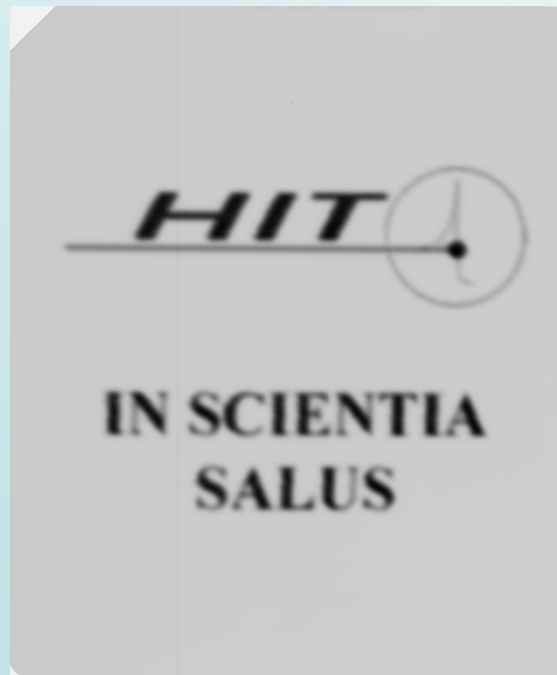
*Optimization of HW and SW for (triggered) image acquisition and data evaluation now under development.*

# Conclusion and Acknowledgements

- There is **no difference between a scientifically used accelerator and a medical accelerator concerning beam diagnostics** equipment, only economic effects may play a role.
- The **speciality of medical accelerators** is the **strong link to the beam consumer**, in this case the **irradiation technology** and the high-conformal **dose application**.
- The common topics “**Instrumentation – Feedbacks – Detector Technology – Quality Assurance**” lead to close collaboration of accelerator and medical physics people in such a facility, resulting in **new and innovative tools for dose application, beam tuning and daily QA**.

# Conclusion and Acknowledgements

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(Intensity modulated  
raster scan,  
 $^{12}\text{C}$  at 430 MeV/u)

***Thank you  
for your  
attention!***