# Introduction to Treatment Planning

#### Alejandro Mazal,

L.DeMarzi, N.Fournier-Bidoz, F.Goudjil, C.Nauraye, S.Delacroix, I.Pasquie, C.Mabit, M.Robilliard, S.Zefkili, M.Auger, A.Patriarca, S.Meyroneinc, C.Devalckenaer, R.Dendale, H.Mamar, V.Calugaru, C.Alapetite, S.Bolle, L.Fevret, S.Helfre, L.Desjardins, A.Fourquet

### **Institut Curie, Paris, France**

Acknowlegments: J-C.Rosenwald, R.Ferrand, T.Lomax, M.Goitein, B.Schaffner, M.Engelsman, N. Schreuder, E.Roelofs, A.Trofimov, J.Flanz, H.Paganetti, H.Kooy, J.Adams, Z.Tochner, E.Hug, H.Giap

Canceropôle, Maastro, France Hadron, Inspira, Saphir, IBA, Varian, Dosisoft, Areva

#### PTCOG-54 San Diego 2015



# Menu of today

- The *process of planning* with protons
- Calculating <u>models</u>
- Planning with <u>passive beams</u>: Compensators and patching
- Planning with <u>scanned beams</u>:

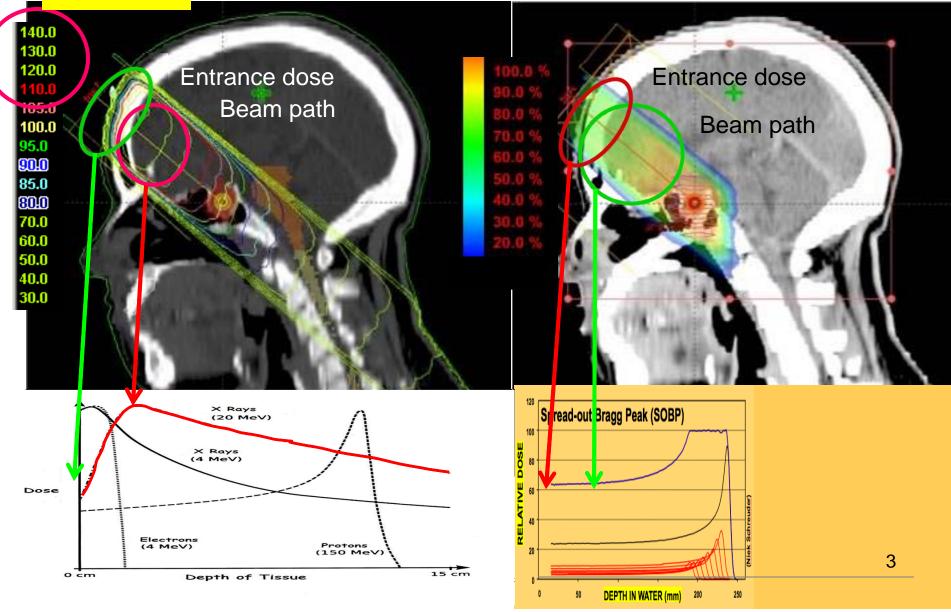
Uniform beams or intensity modulation

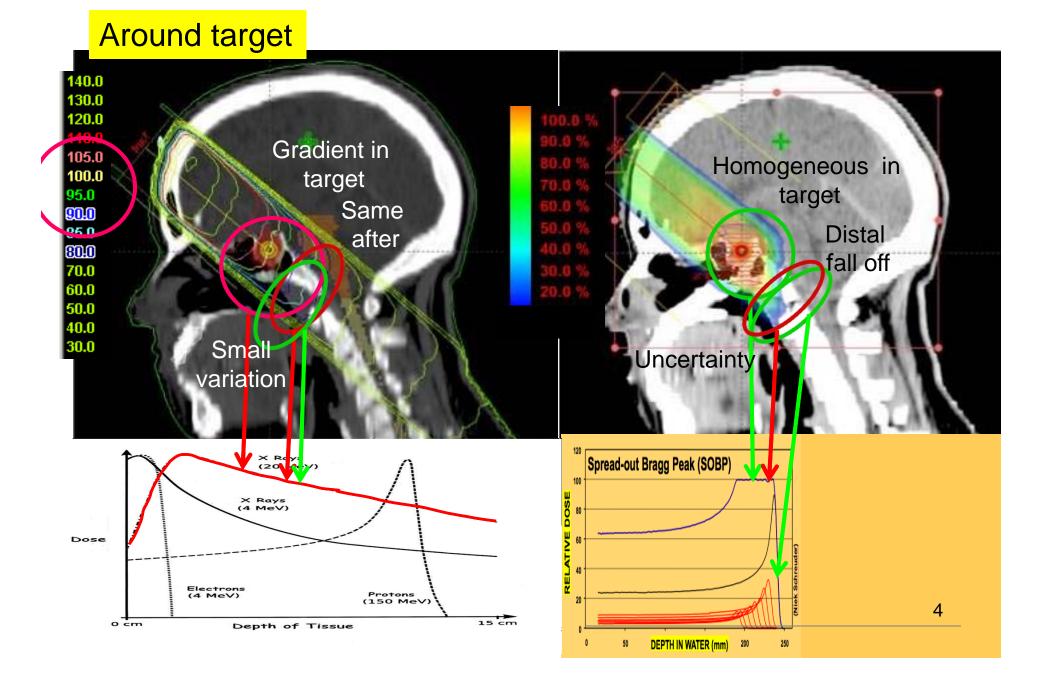
- Organ movements and deformations
- Conclusions
- (lons  $\rightarrow$  see specific talk later)

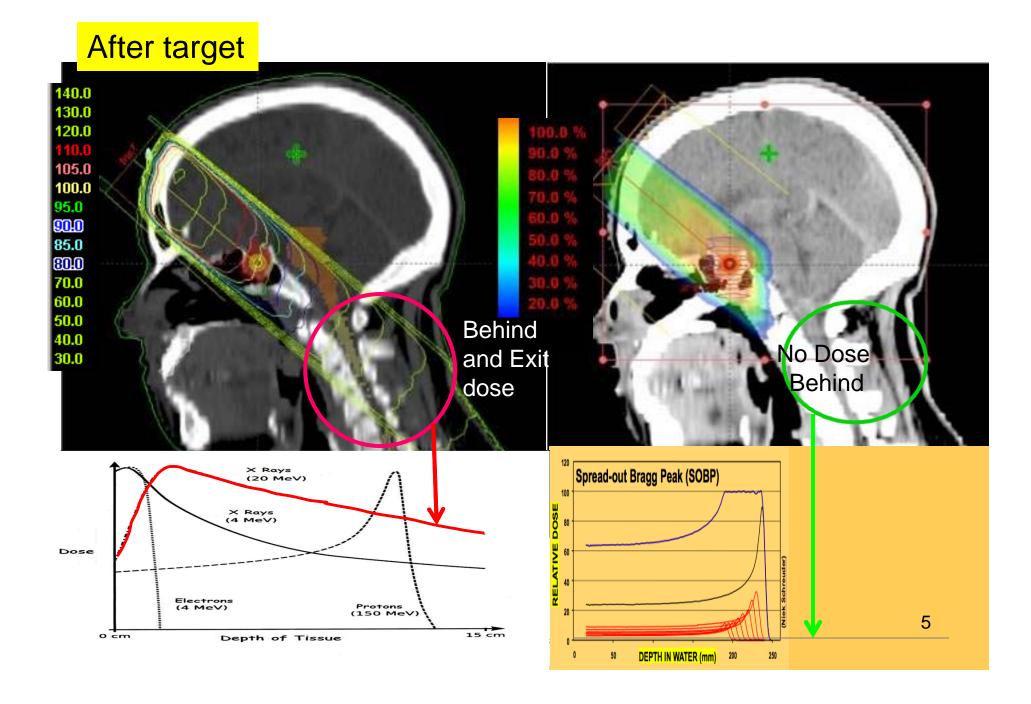


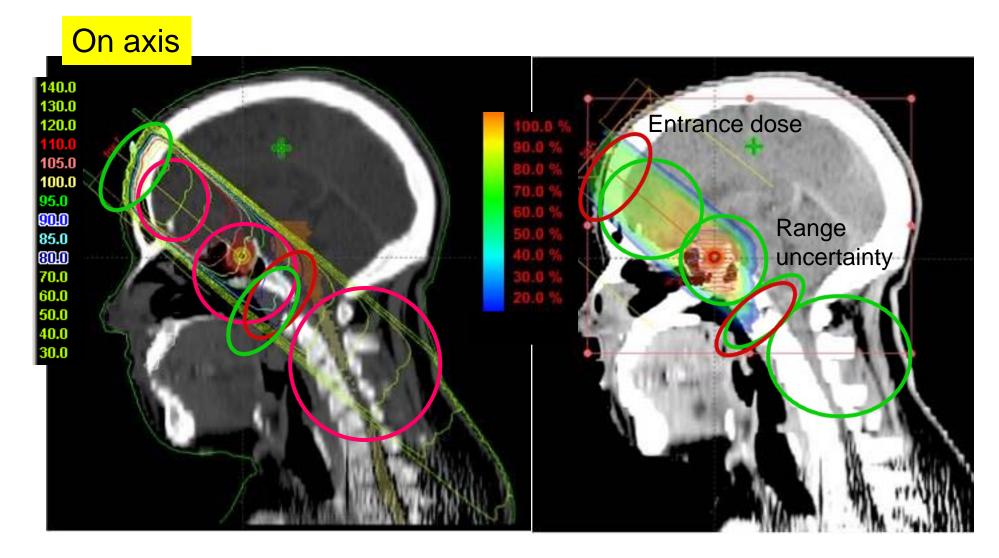
Moving from planning with photons to protons? (Isodoses) (concepts for 1 beam ~ valid for passive and active techniques...)

Entrance

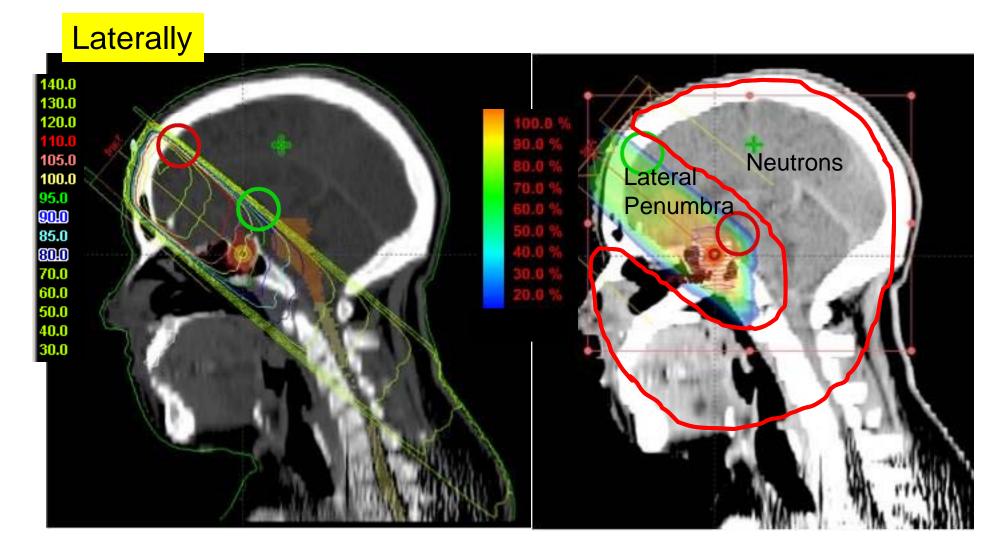




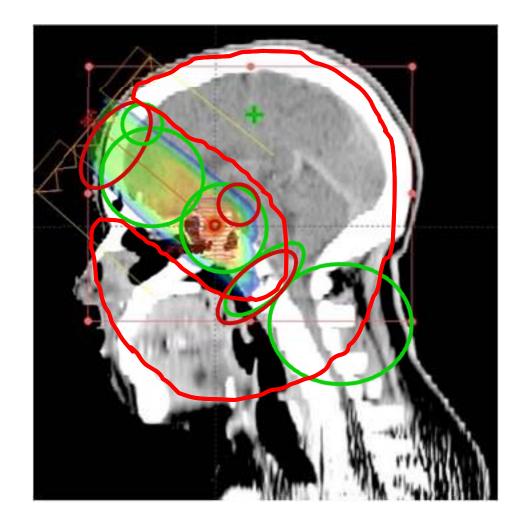














(Sub)liminal message

### **BUT PLANNING**

# **IS NOT ONLY**

### **ISODOSES** and **HISTOGRAMS**

### BUT

## A PROCESS

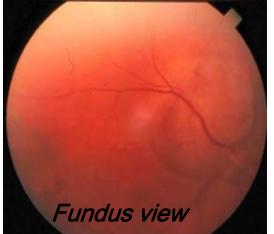


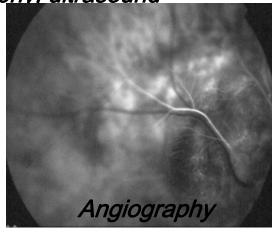
#### **M.Goitein et al**

### <u> The planning process :</u> <u>« First simple case » : Ophthalmologic tumors</u>

### Imaging

*Obtain and inter-register imaging studies : <u>CT. MRI. fundus. angiography. ultrasound</u>* 

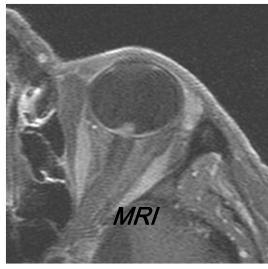


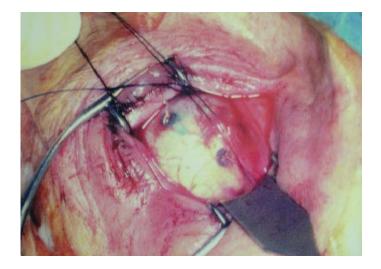


*Immobilisation & reference coordinates* 

*masks, frames,... and/or... use of implanted fiducials* 

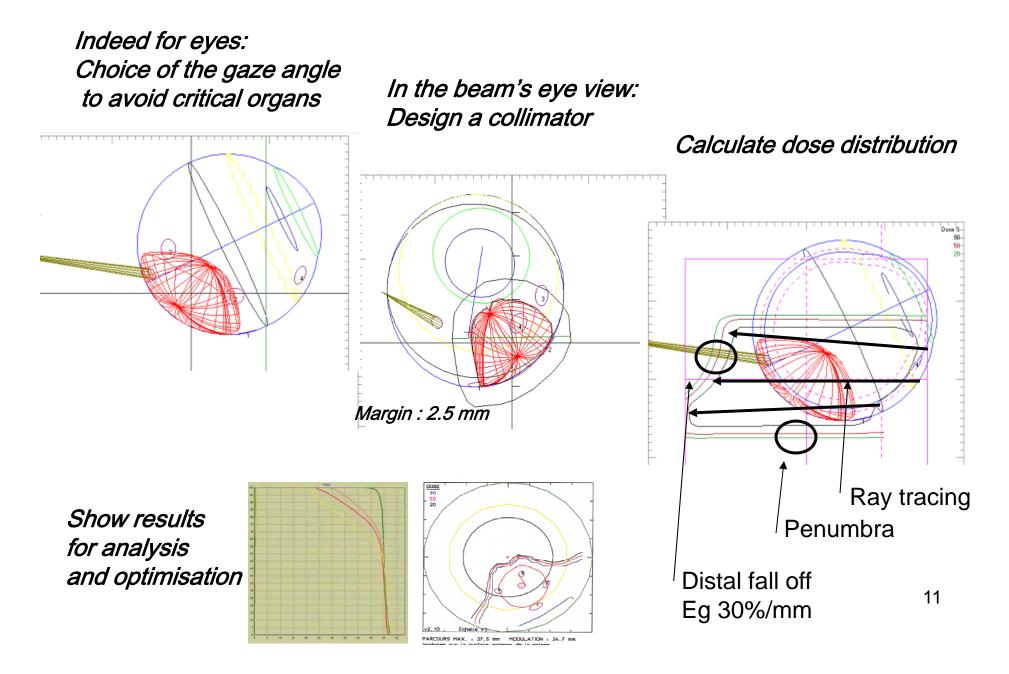




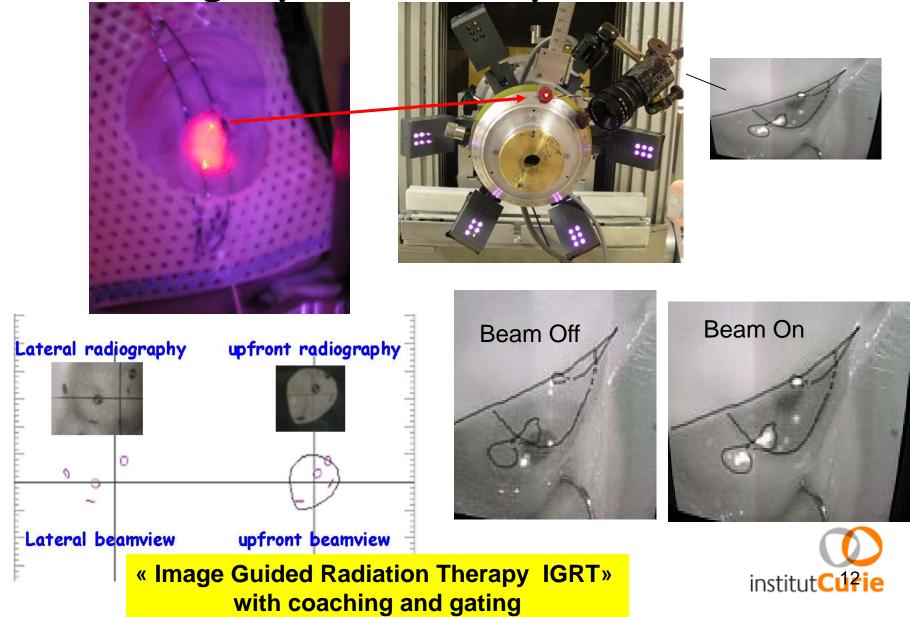




### Delineate target, planning aims and beam design



# Daily set-up control : the Planning System must provide the tools



#### The planning process in general

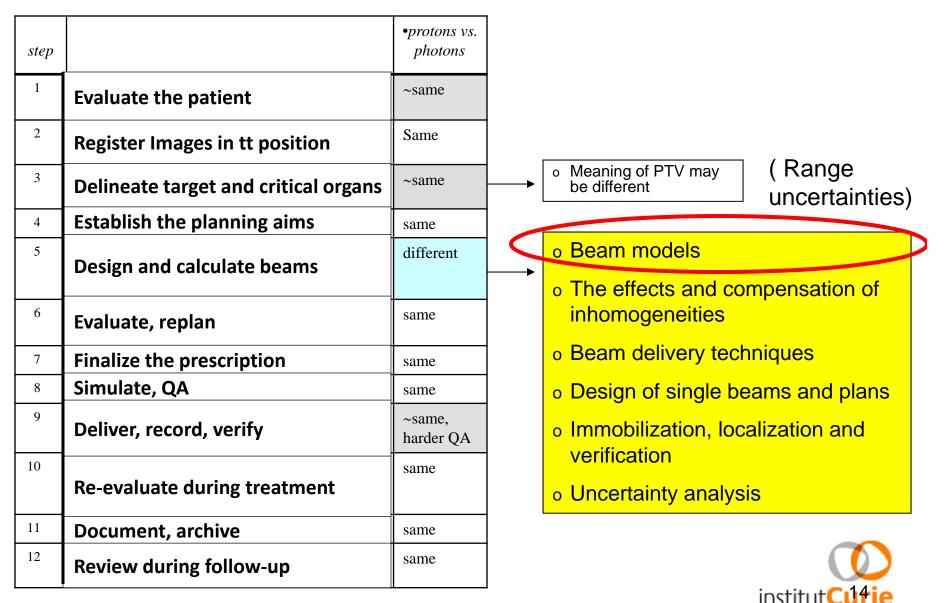
#### (adapted from M.Goitein)

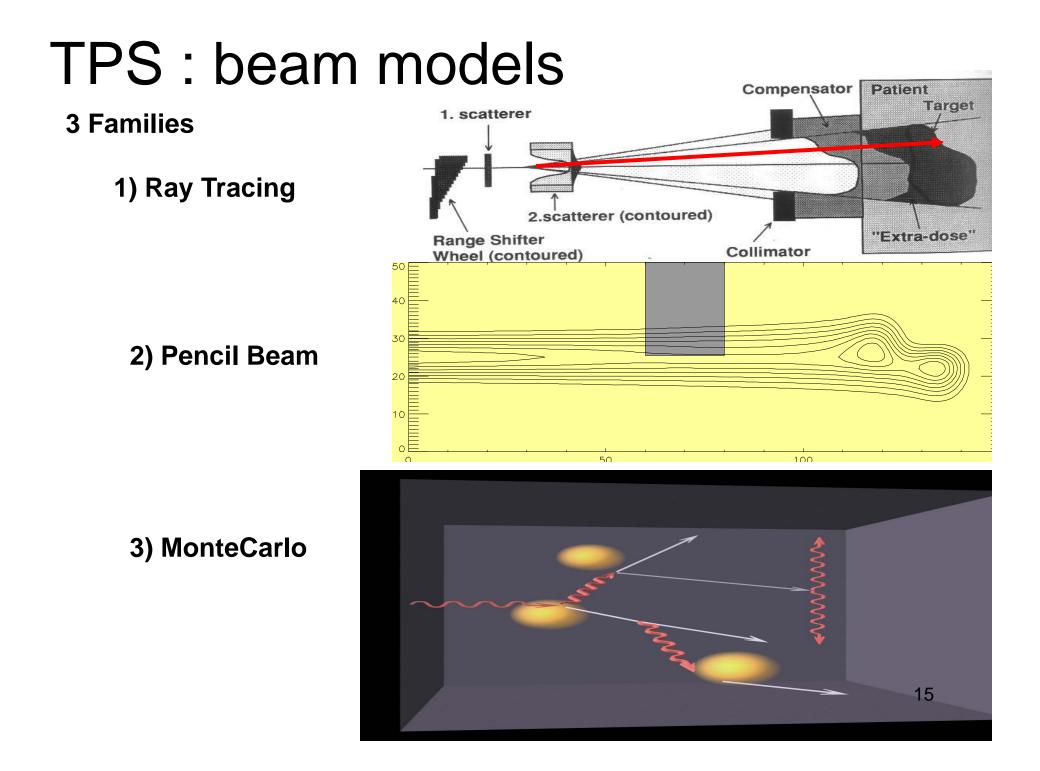
step	
1	Evaluate the patient
2	Register Images in tt position
3	Delineate target and critical organs
4	Establish the planning aims
5	Design and calculate beams
6	Evaluate, replan
7	Finalize the prescription
8	Simulate, QA
9	Deliver, record, verify
10	Re-evaluate during treatment
11	Document, archive
12	Review during follow-up

Steps are common for any approach in RT...



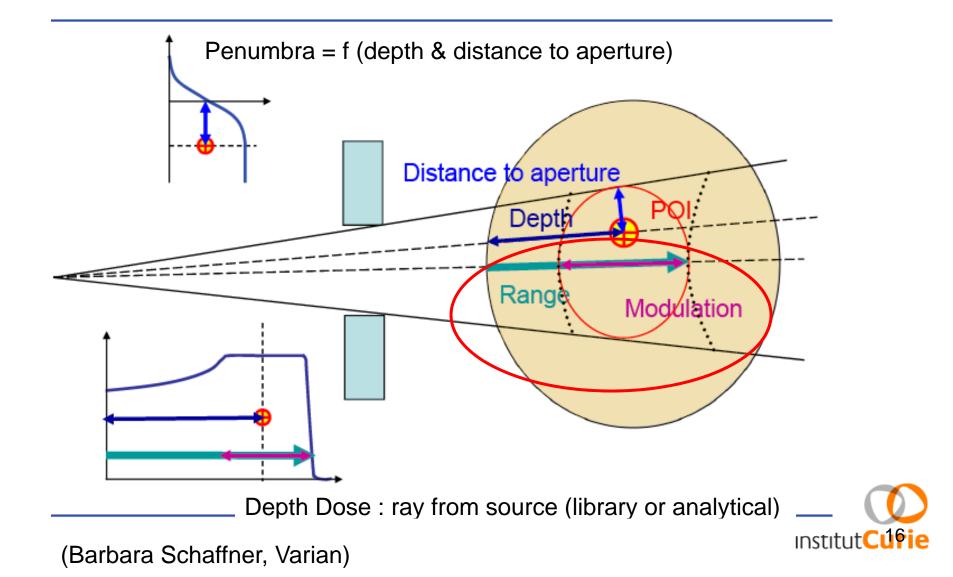
# The planning process in general – and the differences between protons and x-rays





# 1) Ray tracing in passive beams :

### **Broad beam algorithm - Concept**



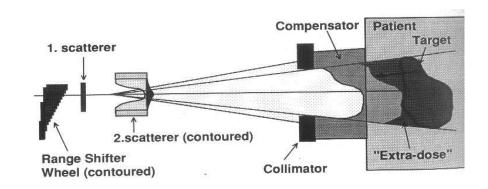
### **Ray tracing :**

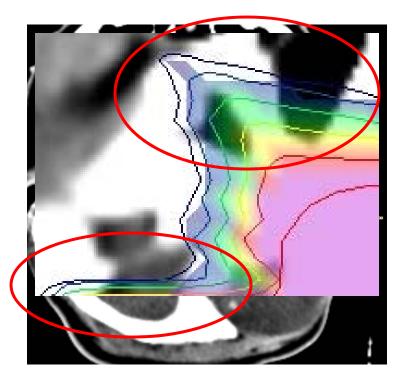
straight protons (no scattering), coming from a (punctual) source

\* lateral penumbra model => takes into account scattering due to :

- initial beam line
- compensator + air-gap
- patient

 $\Rightarrow$  Limitations in inhomogeneous areas and for high gradients in compensators



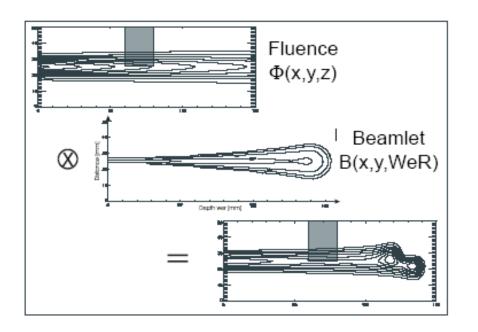


### $\Rightarrow$ Old, simple, fast and relatively efficient



# 2) Pencil Beam

algorithm - Concept



- Principle
  - Convolution of 3D undisturbed proton fluence in air with a 'beamlet' in water.
- In practice
  - Superposition of inhomogeneity corrected beamlets and multiplication with fluence at calculation position.



(Barbara Schaffner, Varian)

### **Pencil Beam :**

• Scattering = broadening of each pencil beam with depth

• Good compromise speed-precision



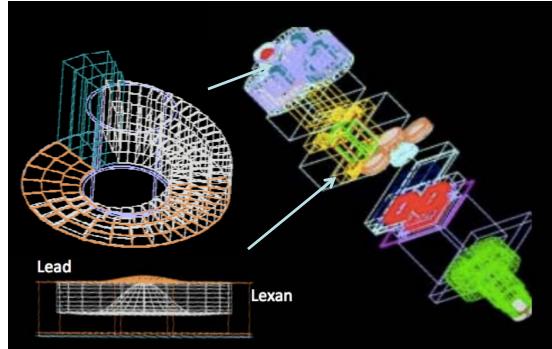
#### The most used at present



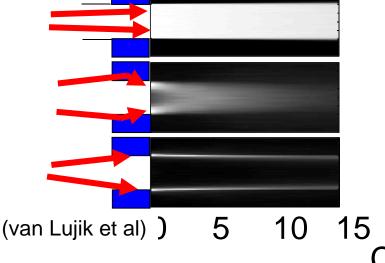
# 3) MonteCarlo

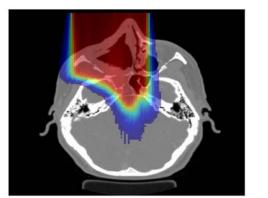
Tracking each particle and all interactions (Geant 4, MCNPX,...):

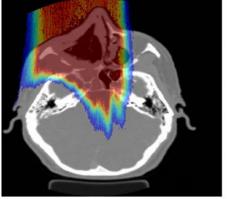
- Beam at the entrance (E,dE,...)
- Treatment Head/nozzle
- 4D if movements
- Patient CT: HU → groups of tissues



Paganetti, Bernardz, et al

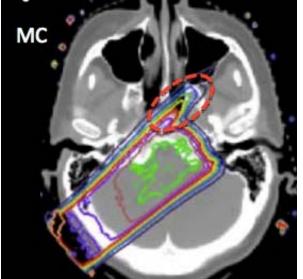






Comparison PB-MC (Paganetti, Trofimov, et al )

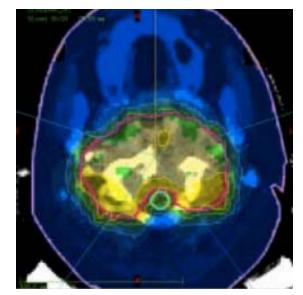
#### **Applications of Monte Carlo :**



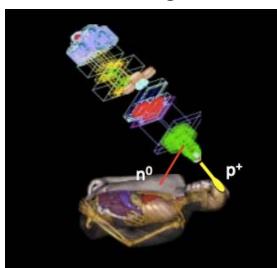
Precise dose calcs with inhomog



Tissue activation for PET QA



Calculation of LET  $\rightarrow$  RBE



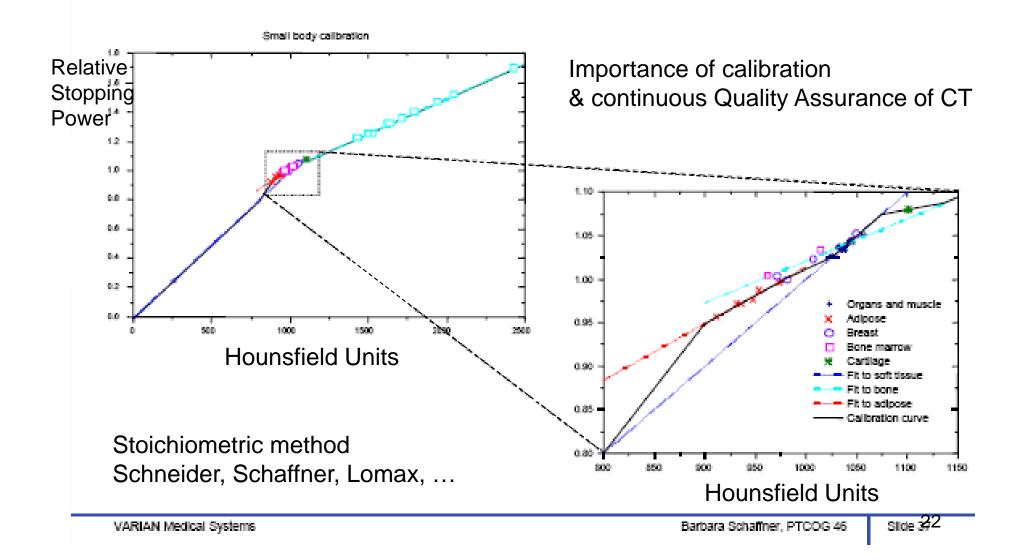
Calculation of neutrons

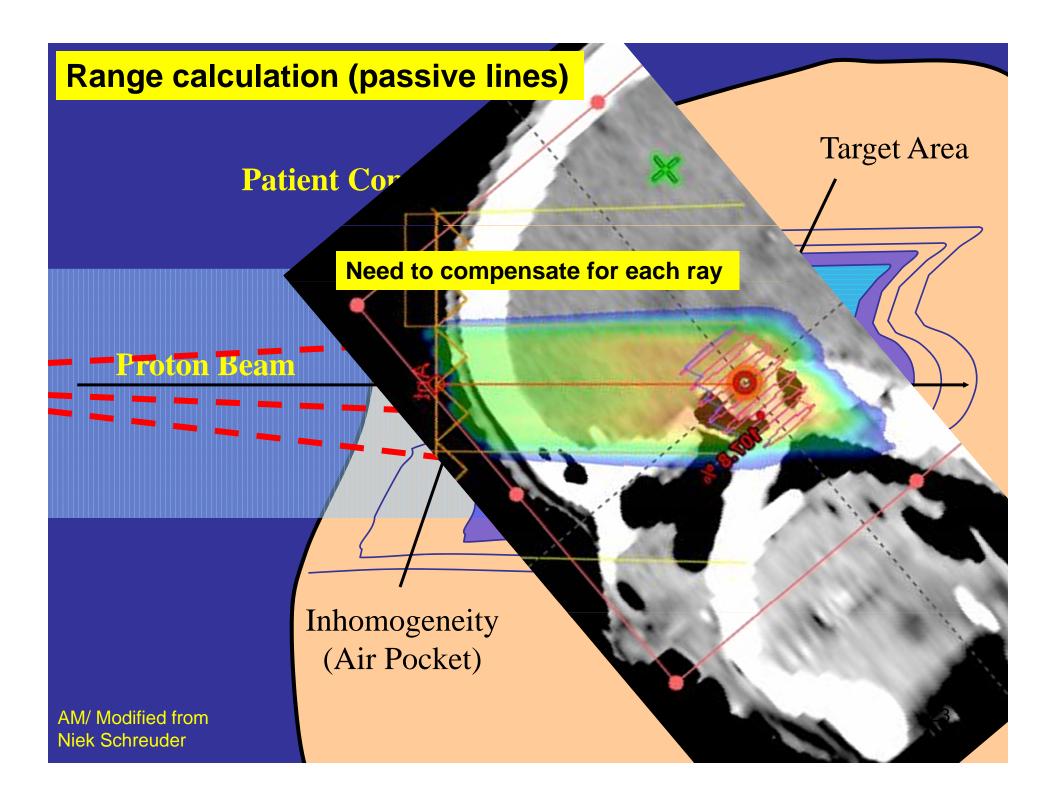
Tor PET GA

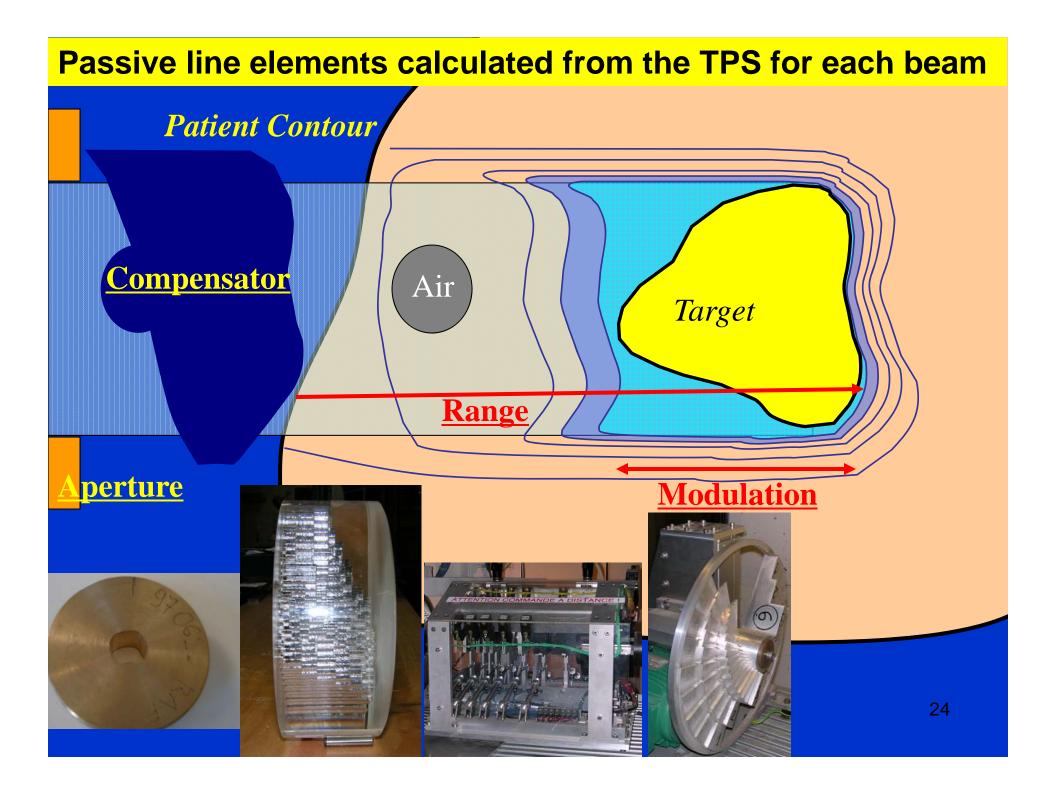
Bednardz, PTCOG49 / (Data from Paganetti, Shin, Espana, Oelfke, Athar, Xu and Bolch)

### 1<sup>st</sup> step:

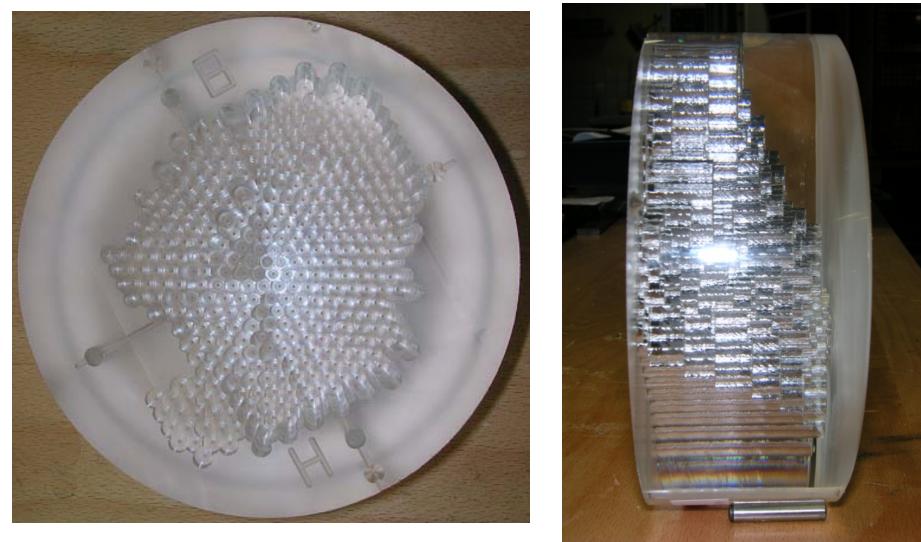
### Conversion from CT Hounsfield Units to Stopping Power (needed for range and dose calculations)



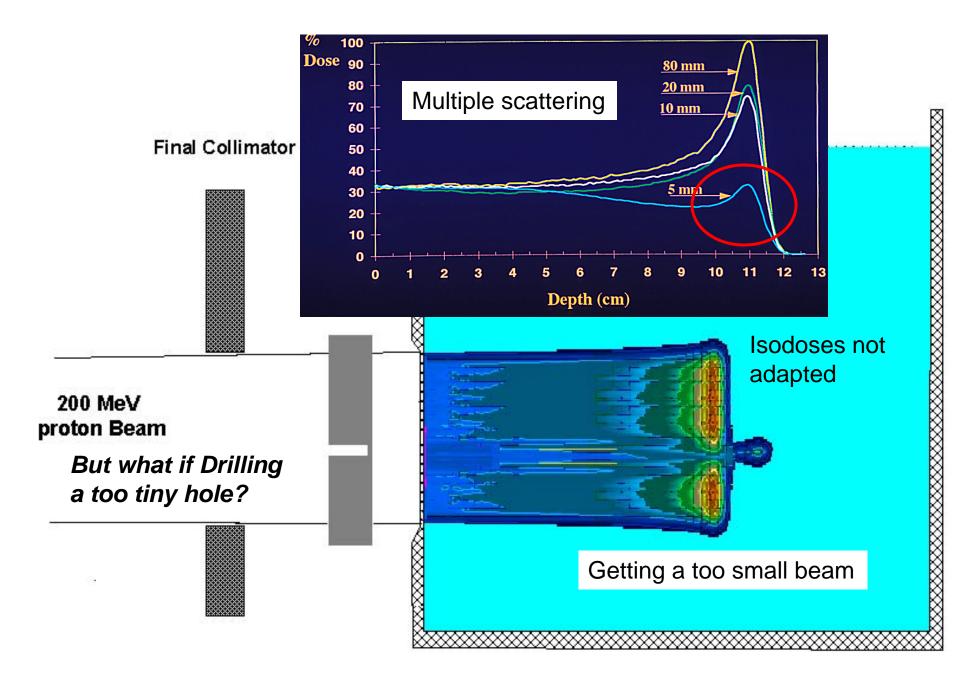


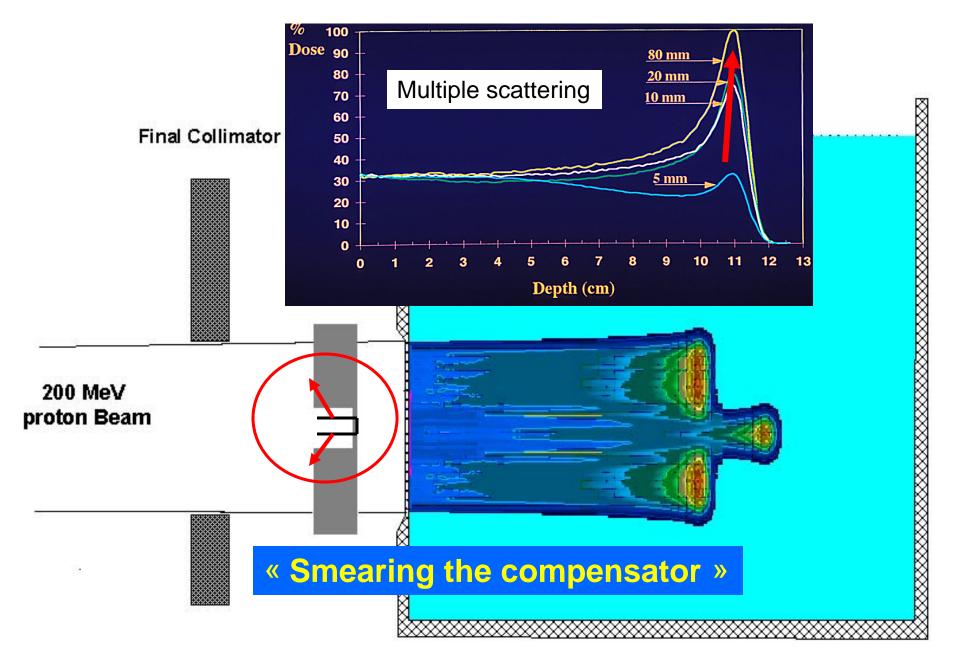


### Example of a compensator



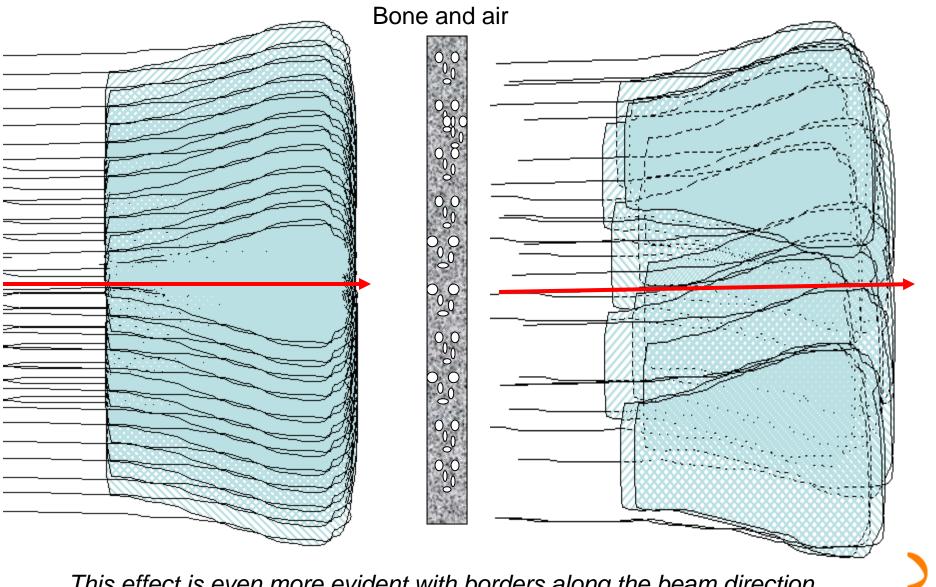




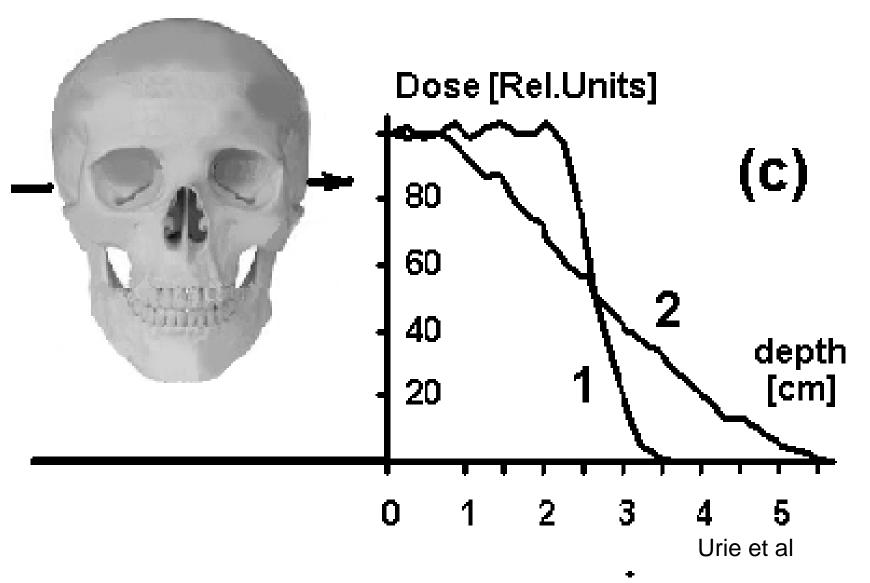


2nd reason to smear : Mis alignements and/or organ movement

If « complex » heterogeneities : multiple scattering effects (all delivery systems)



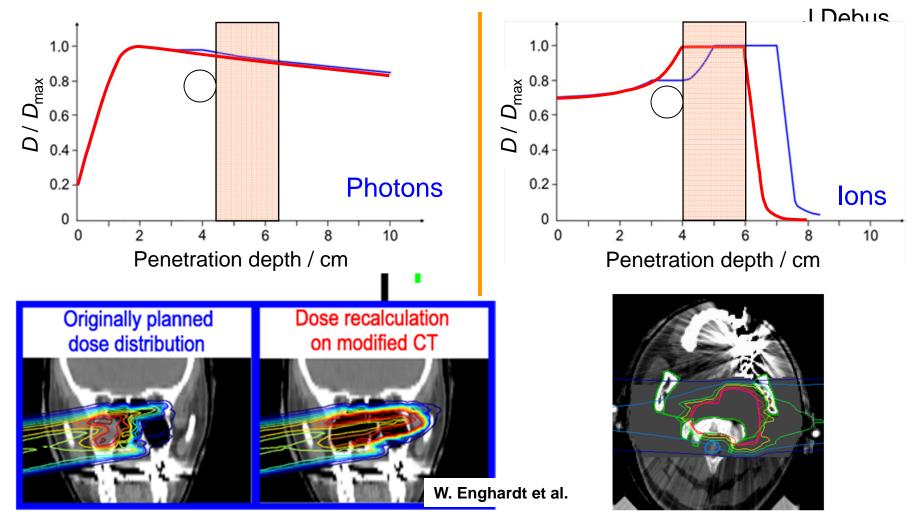
This effect is even more evident with borders along the beam direction... institutC2%ie



- 1) The TPS must calculate that
- 2) Need to change the beam incidence !!



#### Effect of density changes (eg : in the target volume or in the beam path)



Similar effects for CT artifacts, contrast, mispositioning or organ movement

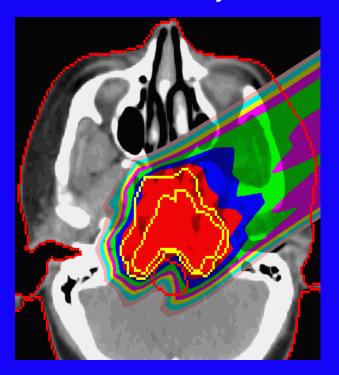
Need to survey the anatomical changes in the path after the planning CT and till the end of the treatment



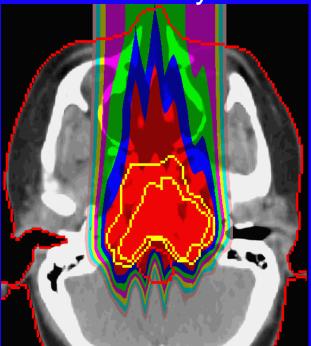
#### Field selection for proton therapy.

Effects on (single field) dose conformity

Example field through relatively homogenous anatomy



Example field through very inhomogenous anatomy



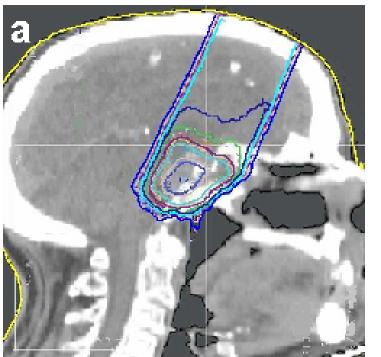


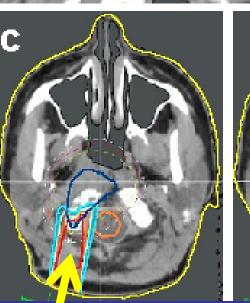
### **<u>Clinical:</u>**

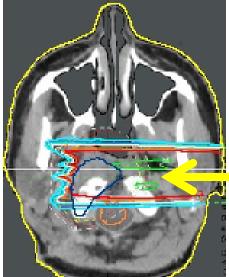
Non coplanar beams

**Photons** + protons

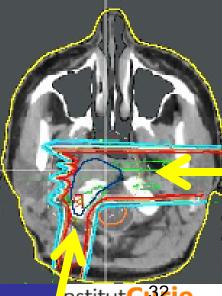
Junctions, "patching"







b

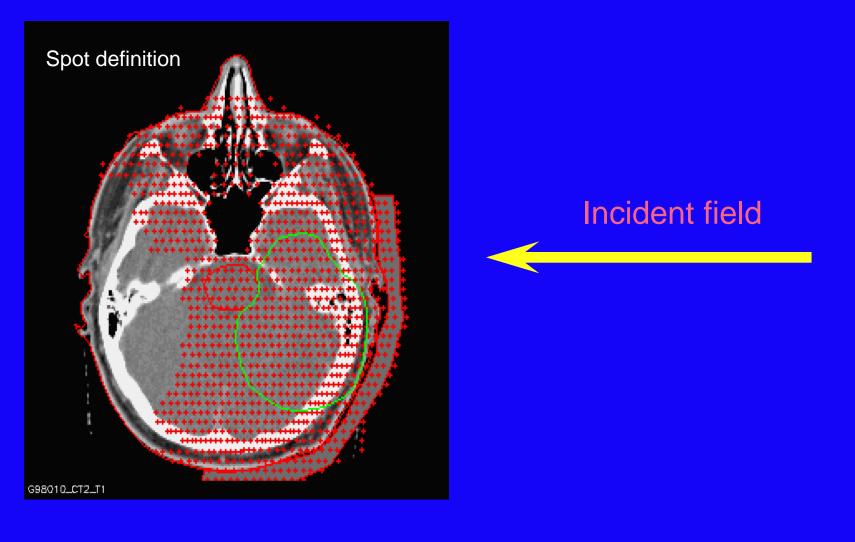


strain :



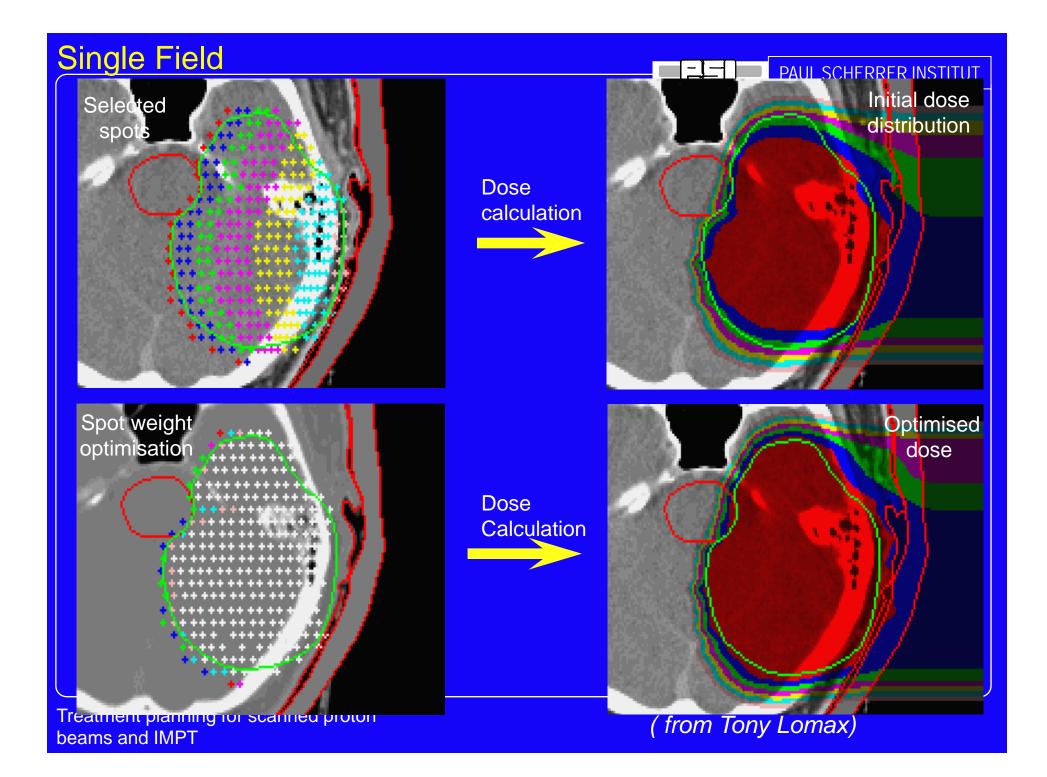


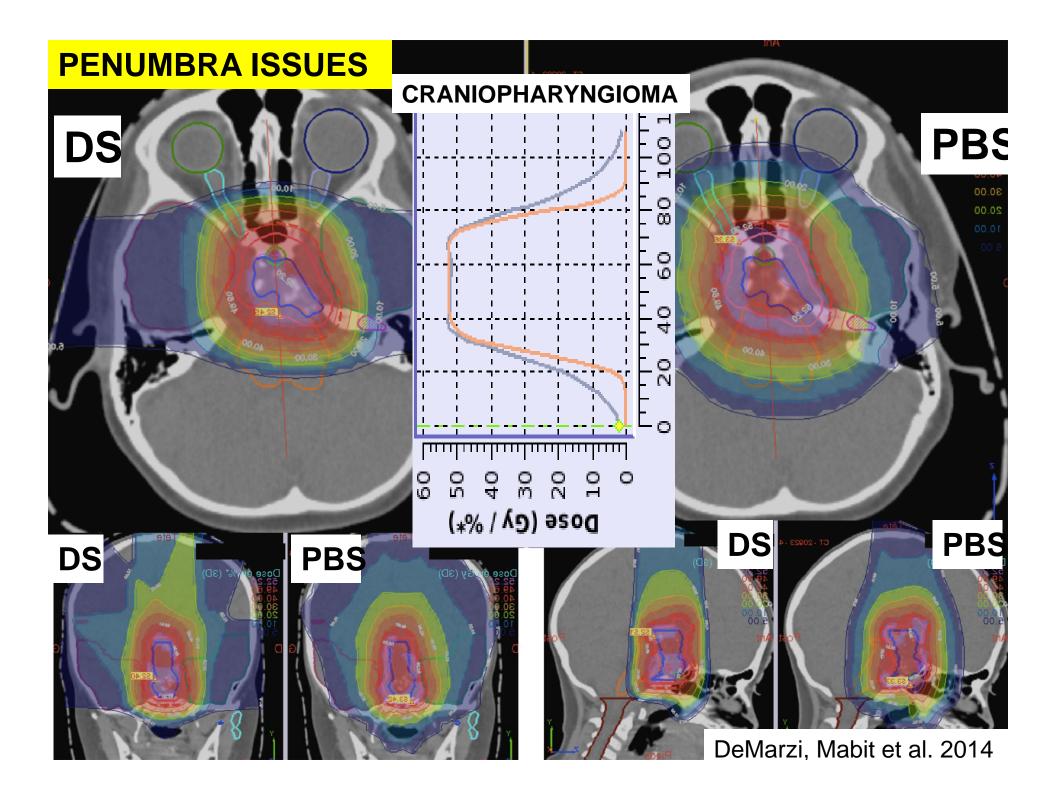
### **Spot Scanning**



Treatment planning for scanned proton beams and IMPT







Ongoing solutions to mitigate :

-Equipment : having smaller spots in the target borders

-Software : **optimizing** spots positions and weights

-Users : adding an aperture for low energy beams

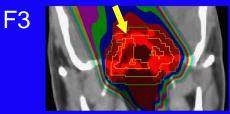
-...

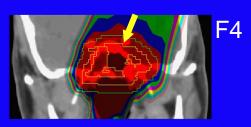
(And have a look to Safai, Bortfeld and Engelsman – Phys. Med. Biol. 53 (2008) 1729–1750)



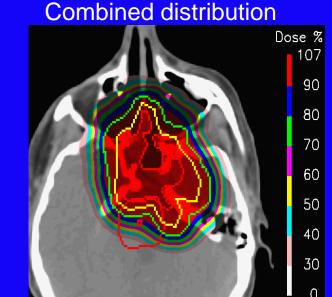
#### Single Field Uniform Dose (SFUD)

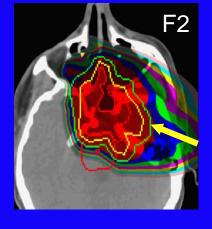
# A SFUD plan consists of the addition of one or more individually optimised fields.





F1





PAUL SCHERRER INSTITUT

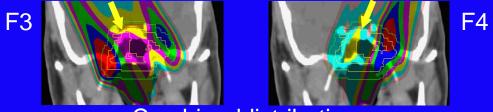
# Note, each individual field is homogenous across the target volume

Treatment planning for scanned proton beams and IMPT

(from Tony Lomax)

Intensity Modulated Proton Therapy (IMPT) 

The simultaneous optimisation of all Bragg peaks from all incident beams



Combineddistribution

122

90

80

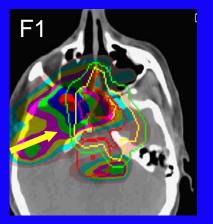
70

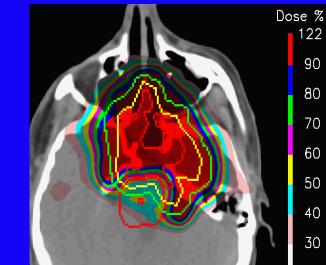
60

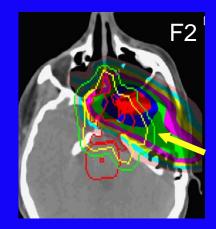
50

40

30







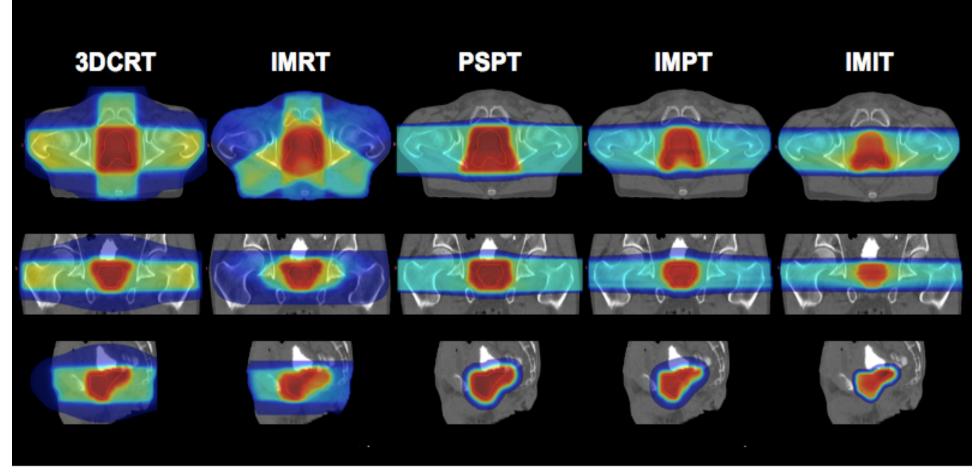
PAUL SCHERRER INSTITUT

Lomax 1999, PMB 44: 185-205

Treatment planning for scanned proton beams and IMPT

(from Tony Lomax)

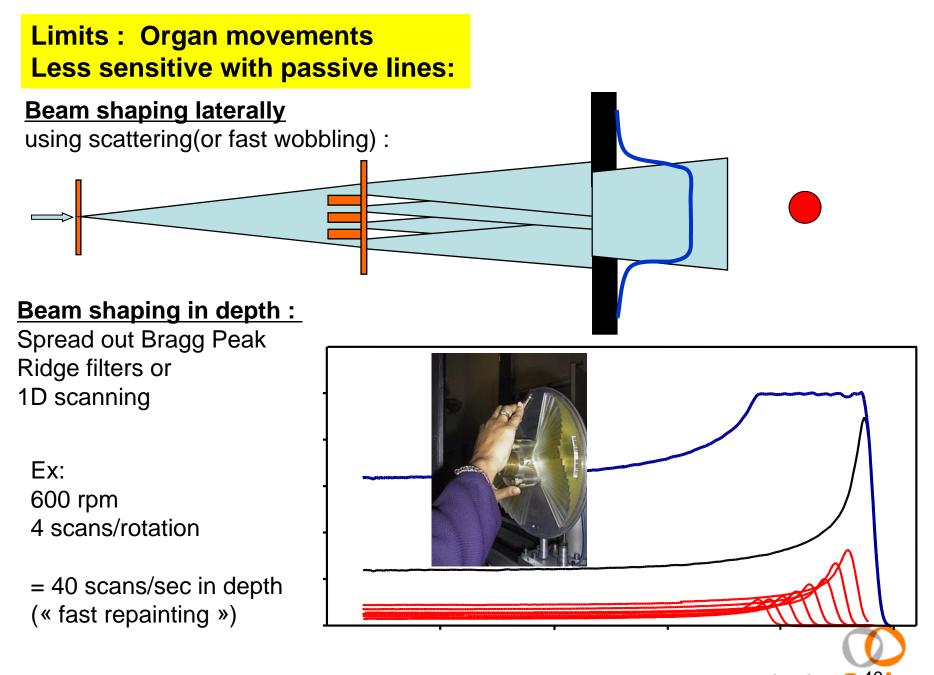
### **Comparative planning**



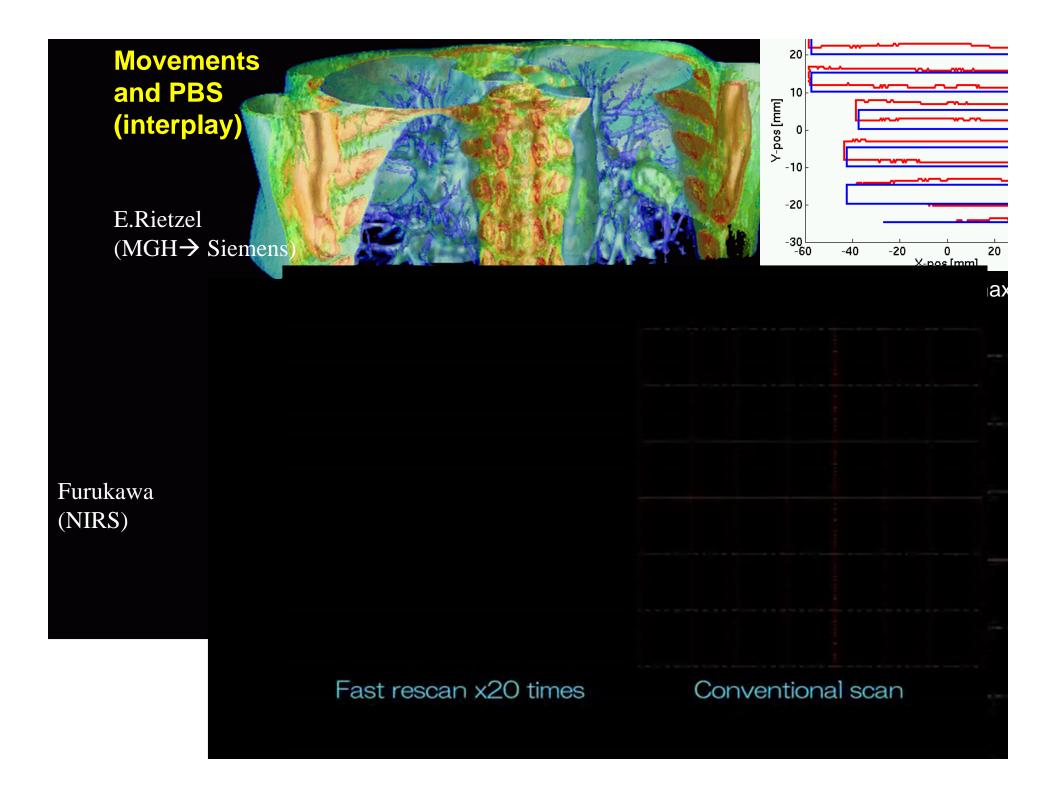
Erik Roelofs et al, ROCOCO Trial, Maastro & 15 institutions involved PTCOG 51, 2011

**Practical examples : See each of the presentations on clinical cases** 





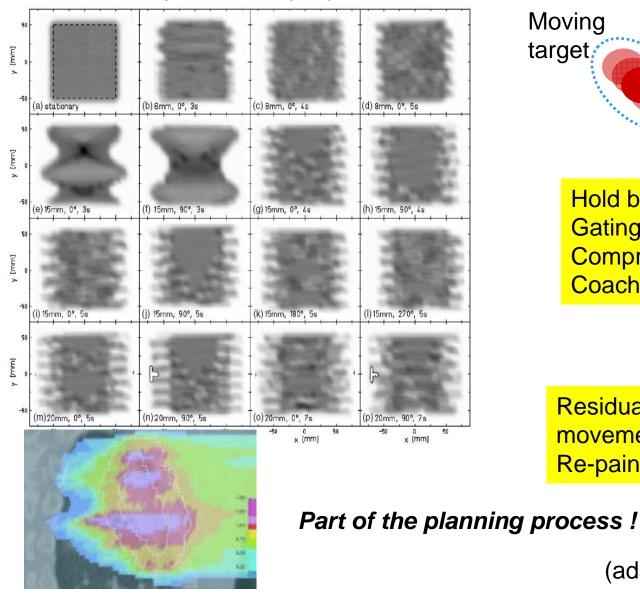
(Less true with the compensator  $\rightarrow$  need to smear the compensator) stitut CdPie

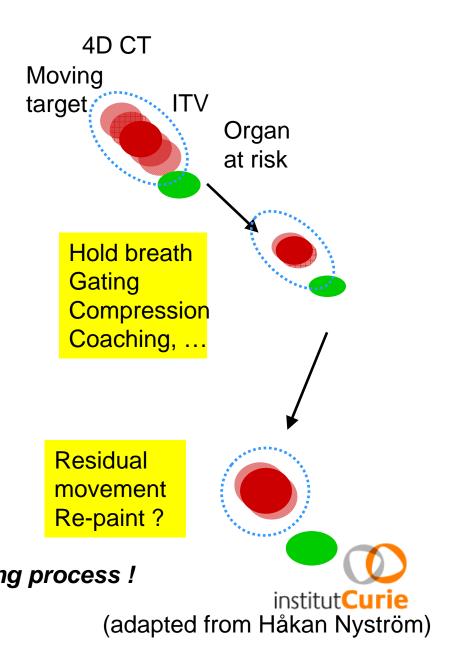


#### PENCIL BEAM SCANNING : very sensitive to Organ Motion

#### Quantification of interplay effects of scanned particle

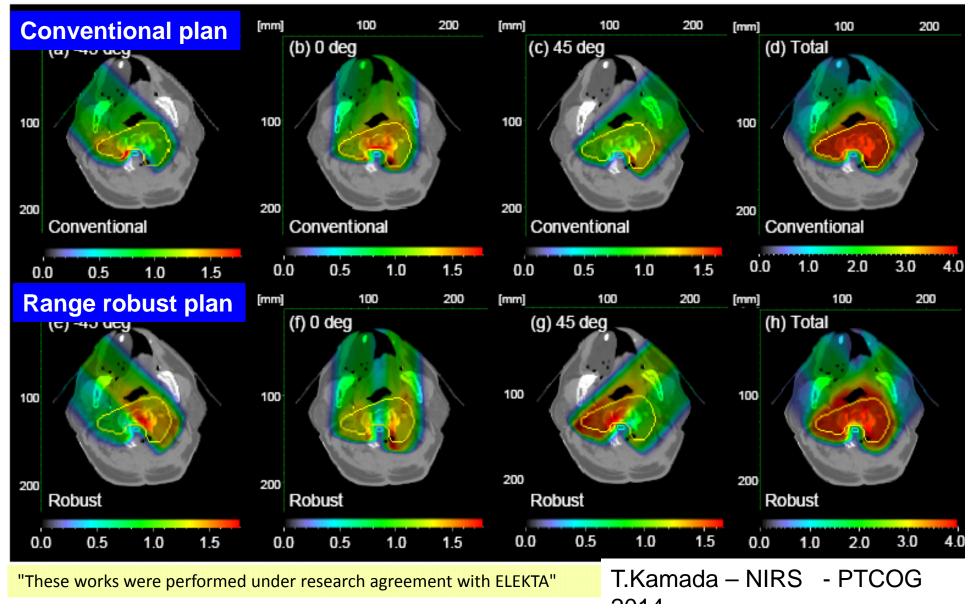
beams and moving targets Christoph Bert, Sven O Grözinger and Eike Rietzel, GSI, Darmstadt, Phys. Med. Biol. 53 (2008) 2253–2265

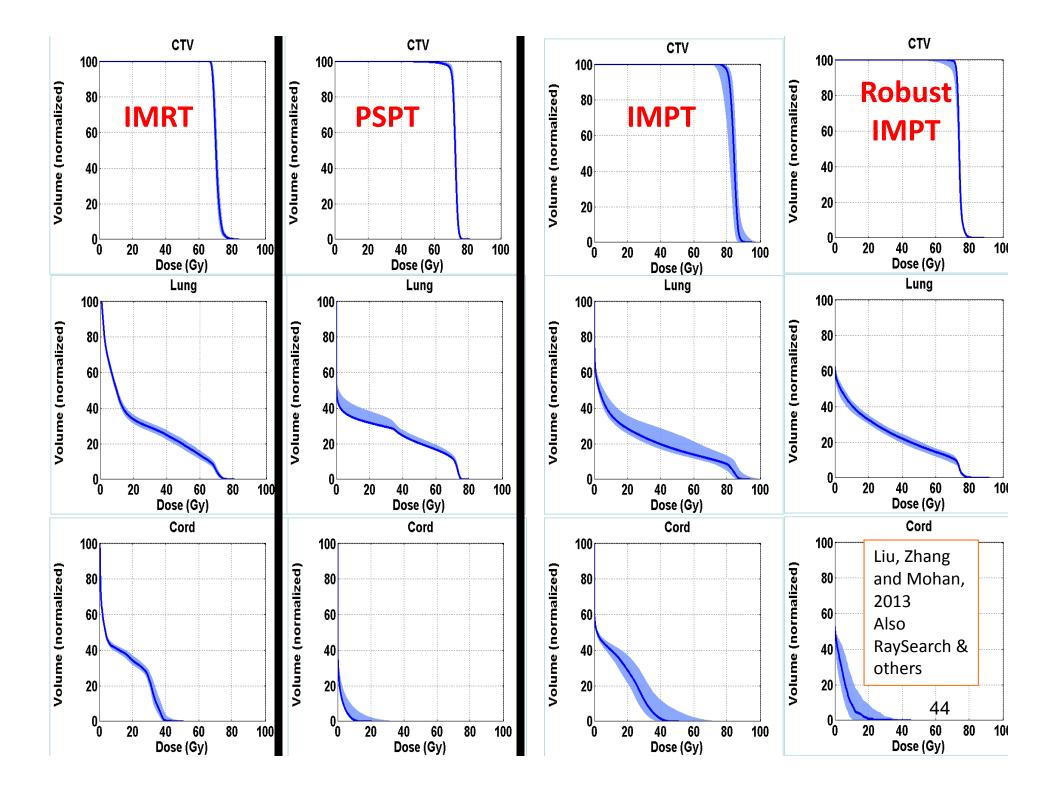




# Lower risk from uncertainties using PBS?

### **Range Robust IMCT for Tumor Surrounding Spinal Cord**





### **Conclusions (I)**

1.Planning with protons is "easy" :

- Fast conformation and reduction in integral dose.
- 2. Different models: Ray tracing, Pencil Beam, Montecarlo,... :
  - Importance of TPS validation, QA and users' experience
- 3. There are limitations :
  - -Entrance dose : multiply beams, combine with photons
  - -Uncertainties in range :

### avoid risky incidences and distal Organ at risk

-Penumbra issues

-Sensitivity to movements, more with dynamic beams :

gating, repainting



### Conclusions (II)

4. Work in synergy with photons, and optimise throughput :

- Fast tools and algorithms
- Need Gantries to plan all incidences as with photons
- 5. Comparative results show in general that :
  - Passive protons ~ > IMXT ( \_\_\_\_\_ integral dose)
  - Intensity Mod PT > IMXT
- 6. TPS evolution towards :
  - MonteCarlo
  - Biological Modeling, mainly for ions
  - Fast and Robust IMPT
  - Adaptive therapy

(need better IGRT & "in vivo" monitoring !)

### Thank You ! Time for Questions ?



# Moving organs and beam scanning : « interplay » & « repainting » concepts

