Patient Specific QA

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Learning Objectives

- 1. Patient specific quality assurance (QA) for passively scattered beams
- 2. QA challenges for pencil beam scanning (PBS)
- 3. In-vivo dose verification techniques



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What Needs QA



"AMARA" or "ALARA"

As Much/Little As Reasonably Achievable/Acceptable

Not Exactly Contradicting!

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QA Evolution



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QA for Passive Scattering



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Passive Scattering



Treatment plan specifies:

Aperture, compensator, range, mod, output factor

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Treatment Configuration



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Aperture



Physical Verification:
Tolerance < 0.5 mm
Imaging Verification:
Tolerance < 1 mm





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Range Compensator



Range in patient depends on thickness $R_p(x, y) = R_{beam} - T(x, y)$

Check T(x,y) at a selected points



Modern technology? Laser, ultrasound, X-ray transmission,



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Range and Modulation



Two definitions: M90 (90%-90%) and M98 (98%-90%)

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M90 versus M98

- M90 -- historical, M98 -- clinically relevant
- Large uncertainties in M90 for large mod
- M90 value may be larger than range
 → impossible to verify



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Measuring Depth-Dose

Sampling interval $t_2 - t_1 = nT$ N = 2, $\delta t < 1$ ms





Lu, Med. Phys. 33 (7), 2006

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Measuring Depth-Dose

- Multi-Layer Ionization Chamber (MLIC)
- 64 plates with 8 chambers per cm
- Cover 8 cm depth



http://physics.harvard.edu/~gottschalk

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Measuring Output Factor MU chambers Isocenter

Output factor depends on ratio r = (R-M)/M

Kooy et al, PMB 48, 2003

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Measuring Output Factor



With aperture and compensator?

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Field Size Effect

Pencil beam calculation (Xio, CMS, Inc):



Depth Dose

Lateral Profile



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Field Size Effect

Measured output change for small field sizes



Go to poster:

"Field Size Dependence of the Output Factor in Proton Radiotherapy" Juliane Daartz, Martijn Engelsman, Marc Bussiere

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Compensator Effect

deep and narrow



Narrow part equivalent to small field

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Information and Work Flow



Statistics leads to confidence!

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Understanding → Less QA

- Identify and correct system instabilities
- Establish model for output prediction
- Use M98 for SOBP specification

Full prediction of SOBP distribution No more evening field cals!

Go to poster:

"A Complete Predictive Model for SOBP Field Delivery" Martijn Engelsman, Hsiao-Ming Lu, David Herrup, Hanne Kooy

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What to do for PBS?



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Pencil Beam Scanning



Pencil specification:

Particle energy (E), Particle count (N), Spot size (σ), trajectory (magnet settings)

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Delivery Methods

Uniform scanning (wobbling)

 fixed scan paths, beam current constant over each layer, fixed range shift from layer to layer, use aperture and compensator

Spot scanning

 treat one spot at a time, beam off between spots, arbitrary range shift between layers

Dynamic scanning

 Beam non-stop within layer, customized scan paths, customized beam current modulation within layer, repainting

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Getting Started

- Understand system capability
- Analyze potential risks
- Develop acceptance standards
- Develop system QA tasks
- Define patient specific QA accordingly
- Measure, analyze, and repeat!

Remember how much you did for IMRT?

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More than IMRT QA

Each layer has own "fluence map" Standard IMRT QA (output and a 2D distribution) Not enough!



2D check for IMRT

One layer off by 8 mm

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In-Vivo Dose Verification



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Sources of Uncertainty

Planning CT HU conversion to stopping power Artifact due to metallic implants Setup errors Variations in position and posture Compensator-patient misalignment Organ motion Lung, liver, pancreas, etc.

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Point Dose Method

Widely practiced in photon/electron therapy

Detectors: MOSFET TLD Diodes







Locations: Surface Cavity Entrance Exit





DVS, implants with wireless reading

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Photon Fields

Measure at one depth, know doses at all depths



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For Protons? Not So Fast! Full dose at point A, but zero dose at point B!



Also need residual proton range at point A

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A Potential Method for DS Beam



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PET for Dose Verification

- Proton and heavy ion beams cause nuclear fragmentation reactions
- Products include positron emitters $^{11}C (T_{1/2}=20.3 \text{ min}), ^{15}O (T_{1/2}=122 \text{ s})$
- Emitters stay at reaction sites
- Activity related to dose distribution

PET image → Dose distribution?



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Activity for Bragg Peak

Monte-Carlo simulations (FLUKA) for proton and carbon beam



(Parodi and Enghardt, Phys. Med. Biol. 45, 2000)

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Activity for SOBP

Measured in polyethylene (PE) phantom

Depth Distribution

Lateral Distribution



(Nishio et al., Med. Phys. 32, 2005)

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Activity Related to Dose

- But, activity is not equal to dose
- Monte-Carlo (MC) simulations can compute both dose and activity distributions
- Compare simulated and measured activity distribution to confirm beam range, dose

Dose vs MC vs PET



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The Process

- Emitter half lives $T_{1/2}=20.3 \text{ min for }^{11}\text{C}$ $T_{1/2}=122 \text{ s for }^{15}\text{O}$ $(T_{1/2}=110 \text{ min for }^{18}\text{F})$
- In-beam imaging (GSI)
- Post treatment PET/CT within 20 min (MGH)





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The Ultimate Proof



Must go to presentation: "In-vivo Imaging in Particle Therapy" Antje Knopf, 4:15PM, Friday, May 23, 2008

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Thank You



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