

Technology and Infrastructure Requirements of a Proton Therapy Centre

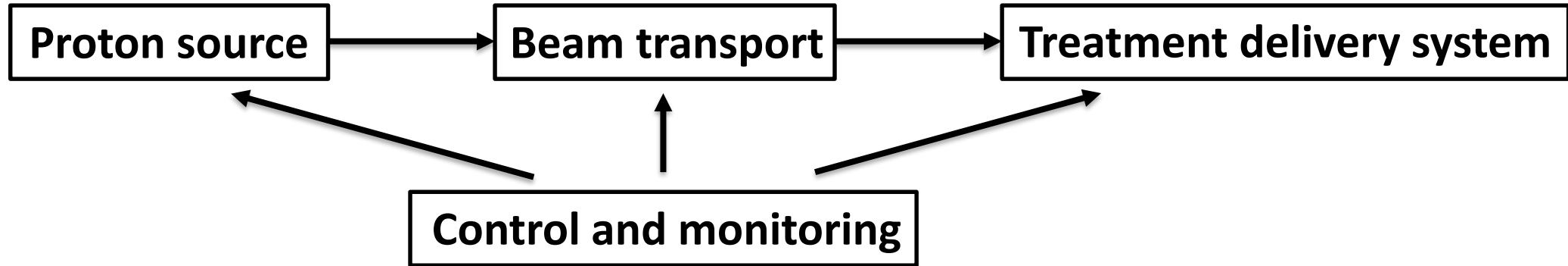
Richard A. Amos, MSc, CPhys, CSci, FIPEM

*Hon. Associate Professor of Proton Therapy
Research Lead for Clinical Proton Therapy Physics
Department of Medical Physics and Biomedical Engineering
University College London*



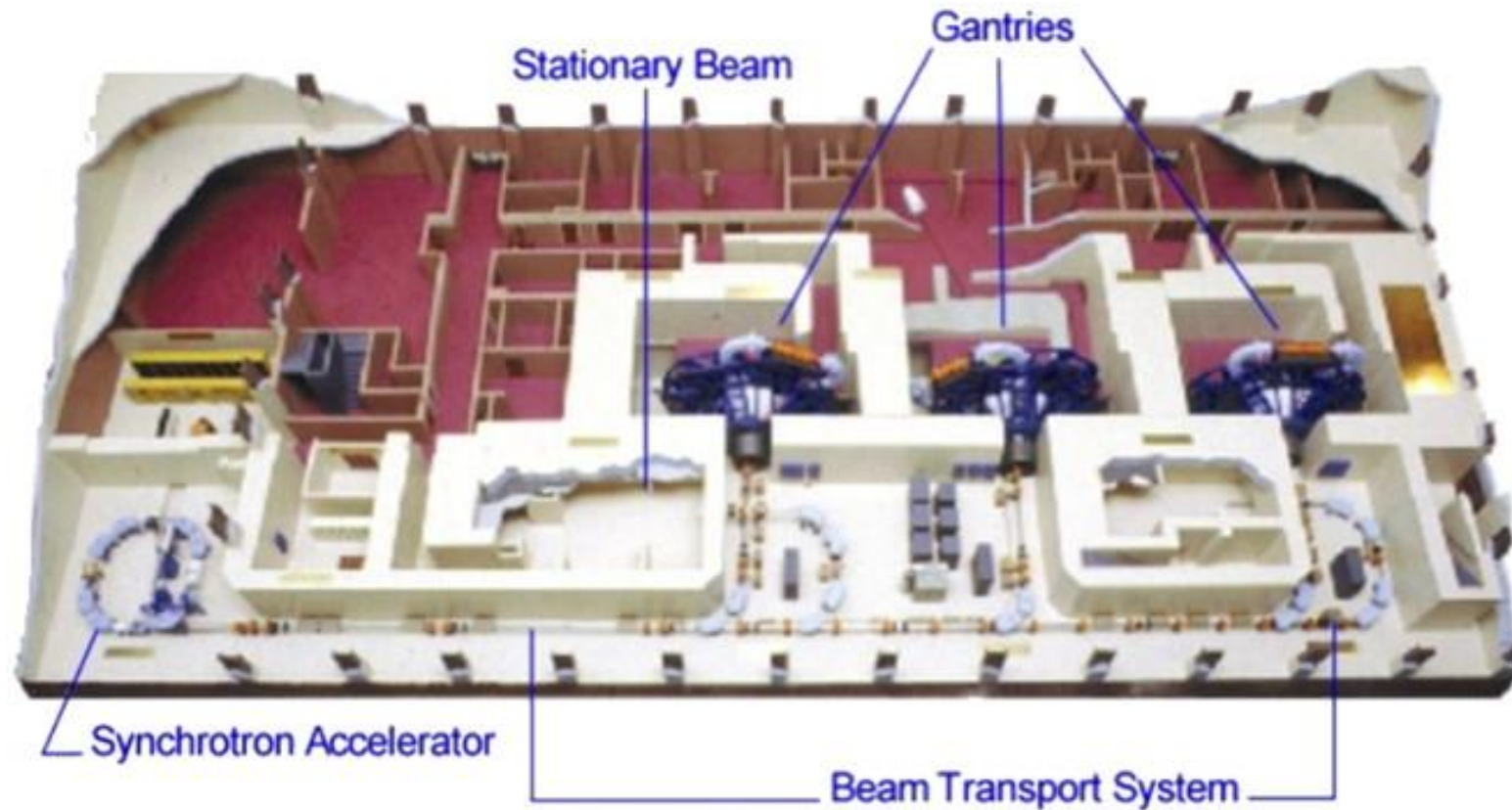
ENLIGHT Annual Meeting
UCL, London, UK
June 25-27, 2018





System specifications defined to meet clinical need:

- Specific needs of case mix
 - Paediatrics
 - Moving targets
 - Complex sites
- Patient throughput requirements
- Capital and maintenance costs
- Additional research capacity

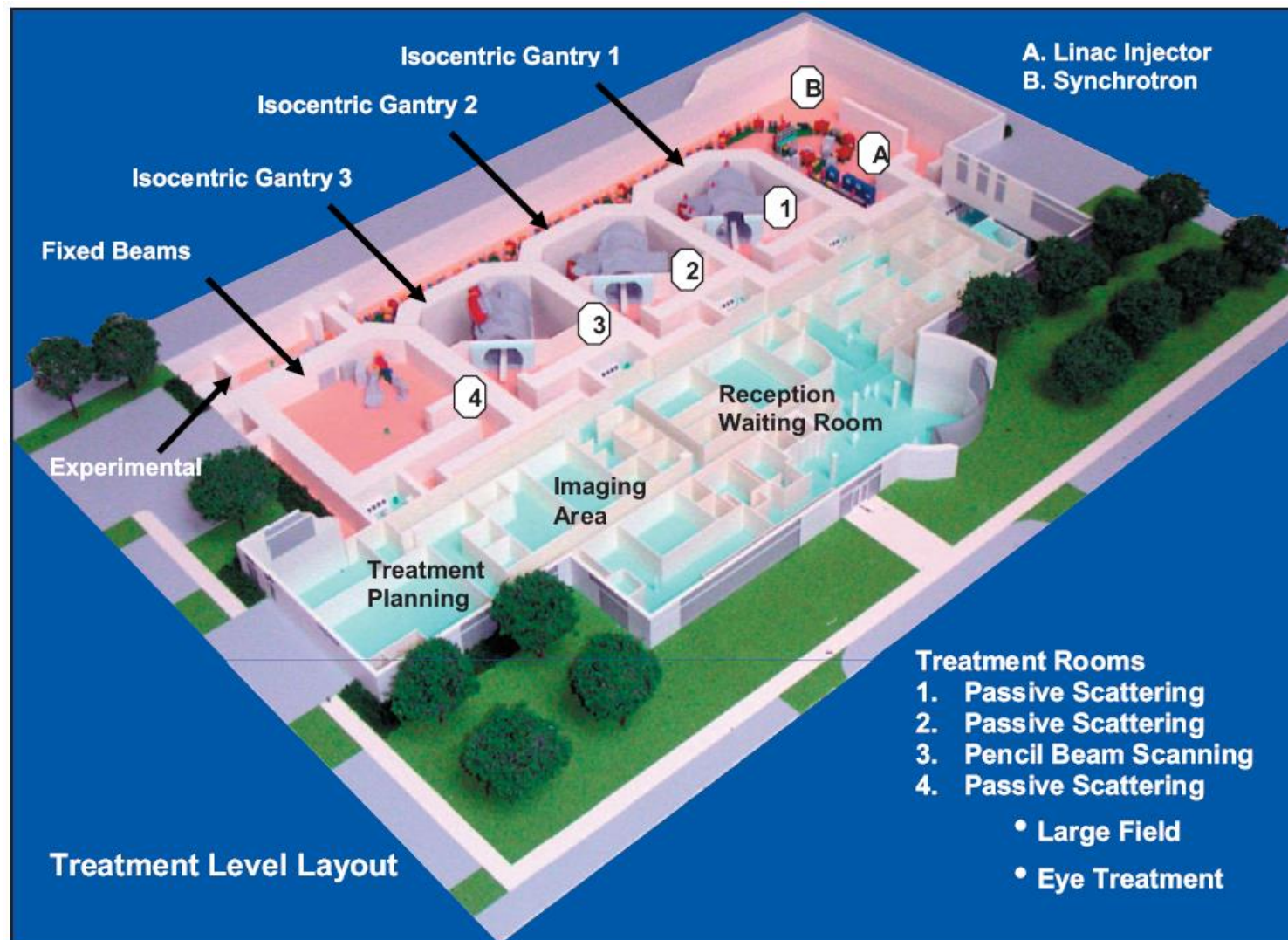


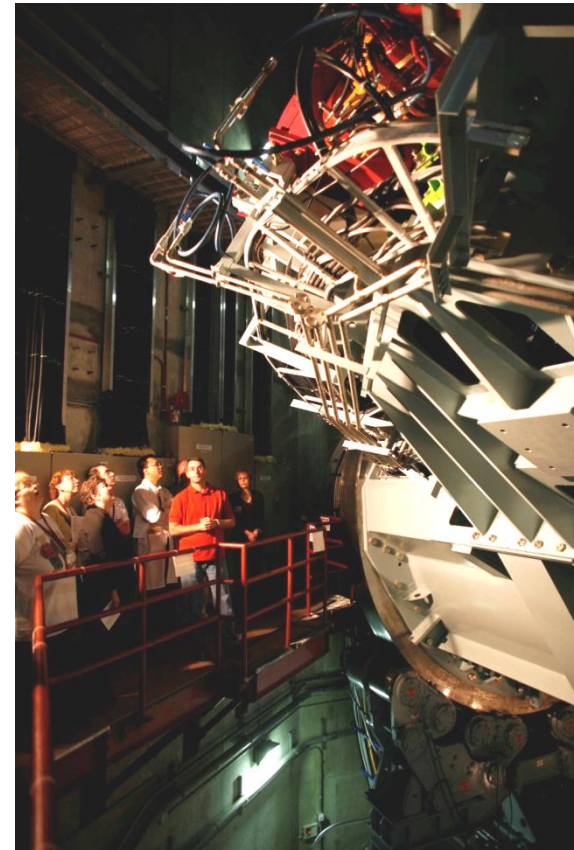
- 250 MeV synchrotron developed in collaboration with Fermi National Accelerator Laboratory
- 3 gantries (passive scattering)
- 1 fixed clinical beamline (passive scattering)
- 1 fixed ocular beamline (passive scattering)
- 1 fixed experimental beamline (passive scattering)

MD Anderson Cancer Center

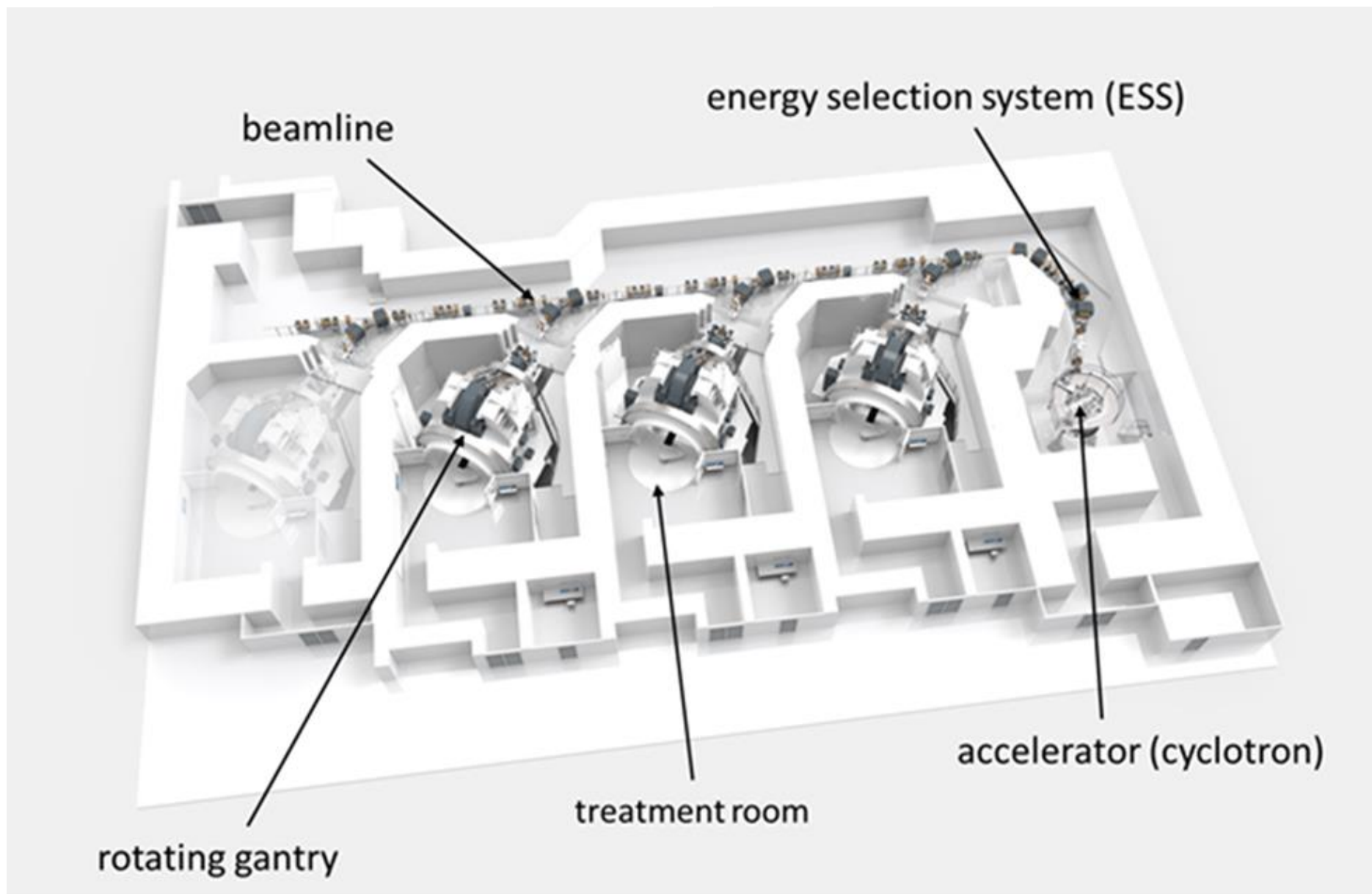
Making Cancer History®

- 250 MeV synchrotron (Hitachi PROBEAT system)
- 3 gantries (2 passive scattering + 1 pencil beam scanning)
- 1 fixed clinical beamline (passive scattering)
- 1 fixed ocular beamline (passive scattering)
- 1 fixed experimental beamline (passive scattering)

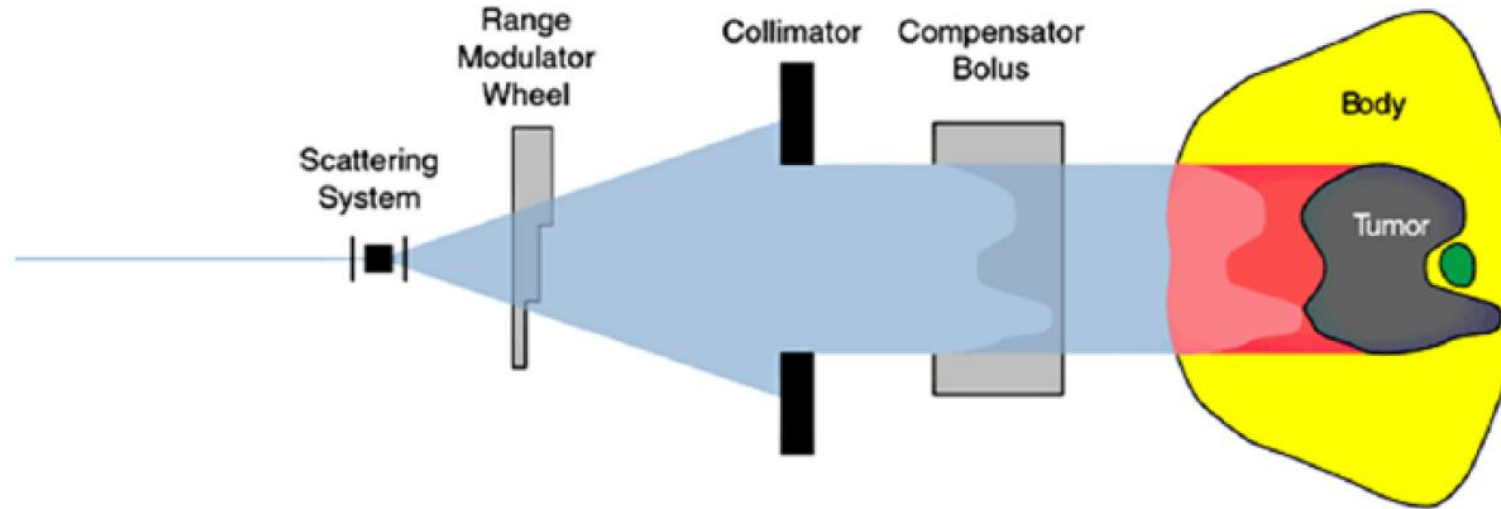




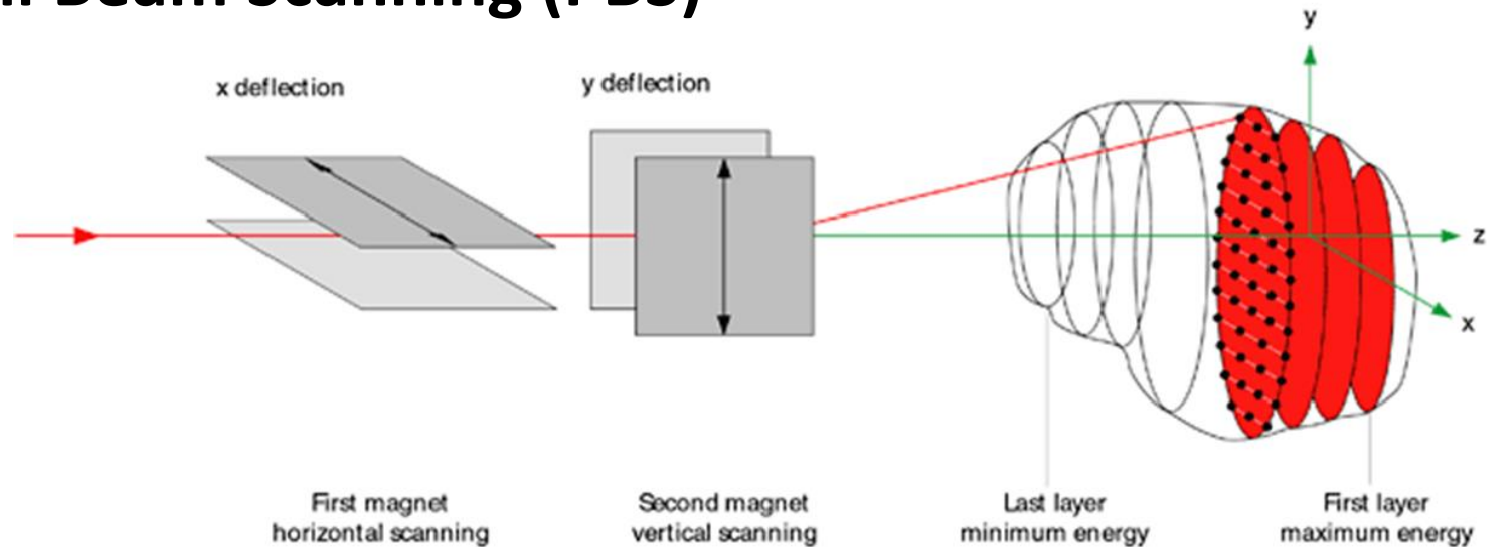
THE UNIVERSITY OF TEXAS
MD Anderson
Cancer Center



Passive Scattering

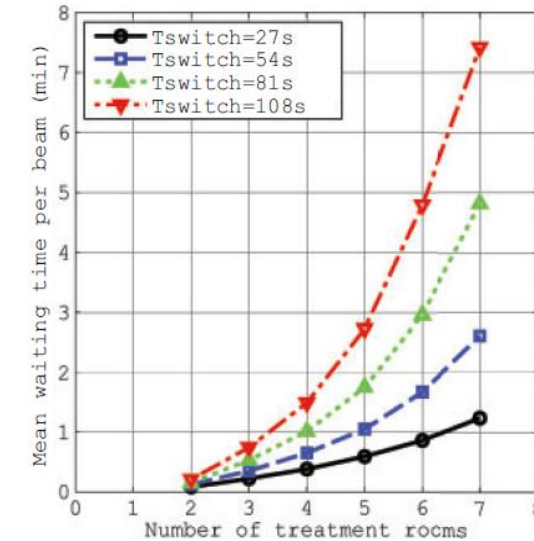
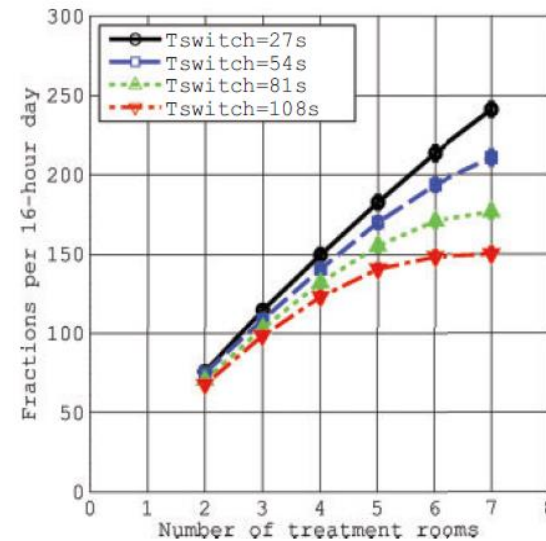
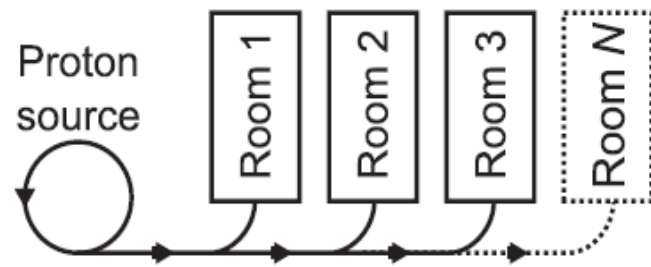


Pencil Beam Scanning (PBS)



Modelling the throughput capacity of a single-accelerator multitreatment room proton therapy centre

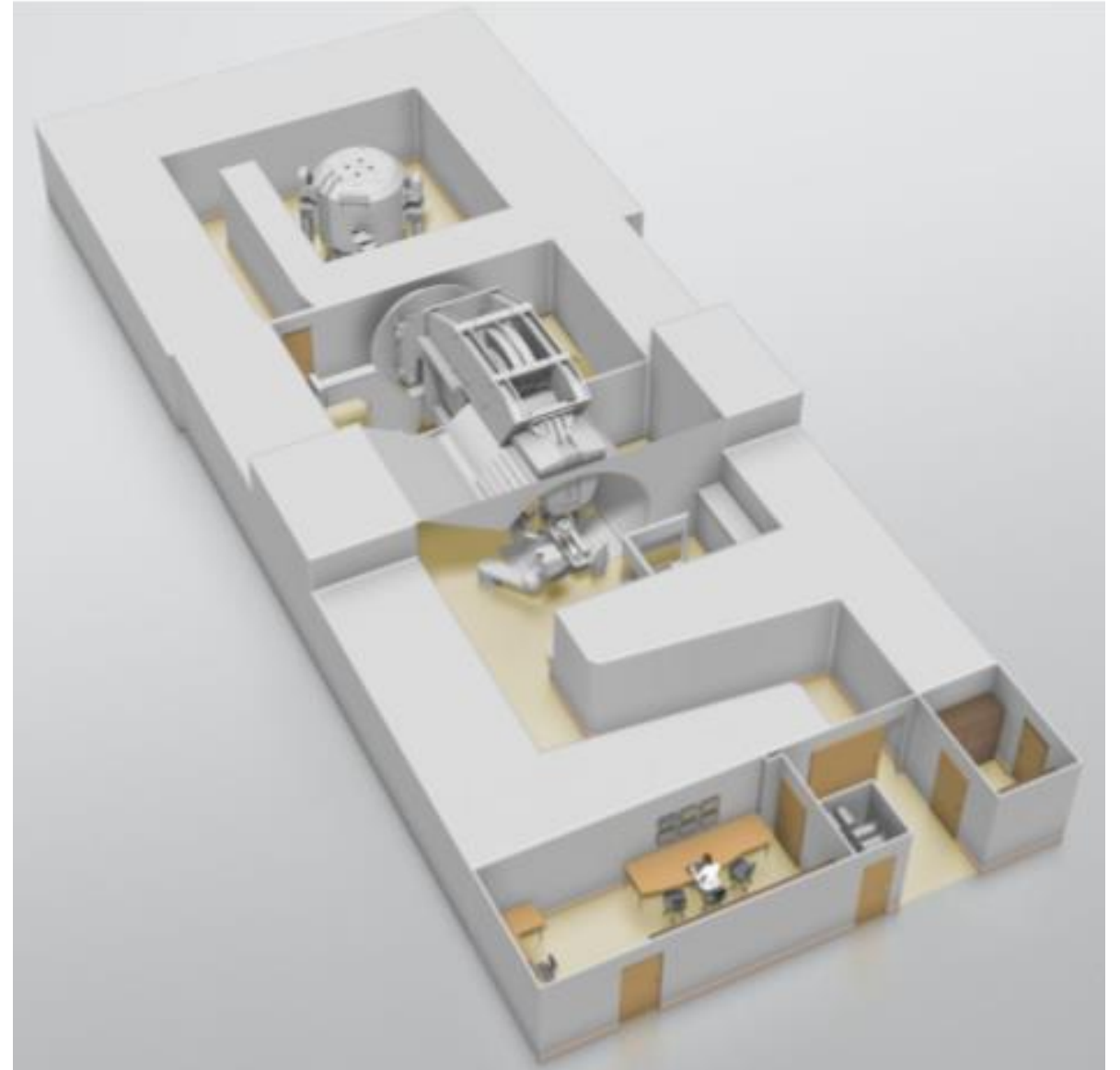
^{1,2}A H AITKENHEAD, PhD, ³D BUGG, BSc, ^{1,2}C G ROWBOTTOM, PhD, ⁴E SMITH, MRCP(UK), FRCR and ^{1,2}R I MACKAY, PhD

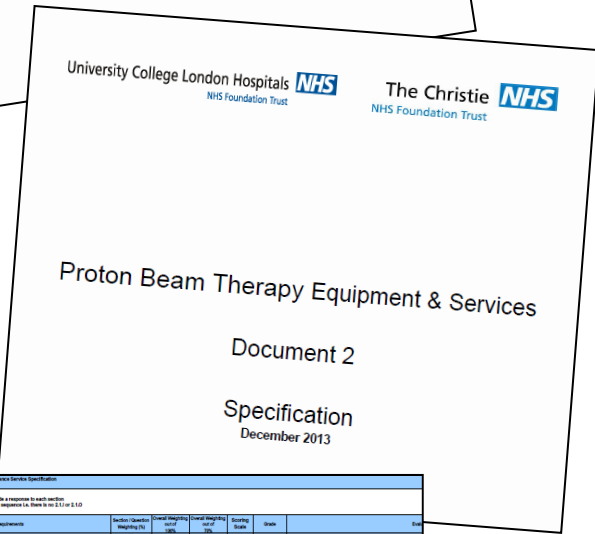


Number of rooms	US caseload			UK caseload		
	t_{fraction} (min)	t_{waiting} (min)	P_{annual} (patients)	t_{fraction} (min)	t_{waiting} (min)	P_{annual} (number of patients)
2	26.6	0.1	546	32.8	0.1	443
3	26.9	0.4	811	33.2	0.4	657
4	27.8	0.7	1046	34.2	0.7	850
5	28.8	1.1	1262	35.5	1.1	1024

Technology evolving:

- Single room facilities
 - Increasing access to wider community
 - Faster construction
 - Efficient
- Partial gantries
 - Space saving
 - Workflow changes
- Accelerator design
 - Higher energy – (*proton radiography*)
 - Cheaper, smaller,....





Question Number	Response Requirements	Section/Question Number/ID	Correct/Incorrect	Weighting/Mark	Answered/Unanswered	Passing/Failed	Mark
1.1A	<p>Question The Bidder must ensure that the Equipment complies with the relevant CE marking, or the Bidder must ensure that the Equipment will achieve CE marking in accordance with the requirements of the relevant CE marking.</p> <p>Response The Bidder must describe the process that will be adopted to achieve CE marking together with the assurance that will be achieved prior to the start of the Equipment. The Bidder must confirm that CE certification will be provided at an additional cost to the Trust. Bidder must describe how any parts of the Equipment that do not or will not CE mark.</p> <p>Options Provide proof of approval evidence or provide written statement. Provide the proposed evidence that CE marking will be achieved in order for treatment to commence by the start of the Equipment.</p>	0%	0%	0%	Pass/Fail	Fail	
1.1B	<p>Question The Bidder must ensure that the control cabinet complies with the relevant CE marking, or the Bidder must ensure that the control cabinet will achieve CE marking in accordance with the requirements of the relevant CE marking.</p> <p>Response The Bidder must describe the process that will be adopted to achieve CE marking together with the assurance that will be achieved prior to the start of the Equipment. The Bidder must confirm that CE certification will be provided at an additional cost to the Trust. Bidder must describe how any parts of the Equipment that do not or will not CE mark.</p> <p>Options Provide proof of approval evidence or provide written statement. Provide the proposed evidence that CE marking will be achieved in order for treatment to commence by the start of the Equipment.</p>	0%	0%	0%	Pass/Fail	Fail	
1.1C	<p>Question The Bidder must ensure that the control cabinet complies with the relevant CE marking, or the Bidder must ensure that the control cabinet will achieve CE marking in accordance with the requirements of the relevant CE marking.</p> <p>Response The Bidder must describe the process that will be adopted to achieve CE marking together with the assurance that will be achieved prior to the start of the Equipment. The Bidder must confirm that CE certification will be provided at an additional cost to the Trust. Bidder must describe how any parts of the Equipment that do not or will not CE mark.</p> <p>Options Provide proof of approval evidence or provide written statement. Provide the proposed evidence that CE marking will be achieved in order for treatment to commence by the start of the Equipment.</p>	0%	0%	0%	Pass/Fail	Fail	
1.1D	<p>Question The Bidder must ensure that the control cabinet complies with the relevant CE marking, or the Bidder must ensure that the control cabinet will achieve CE marking in accordance with the requirements of the relevant CE marking.</p> <p>Response The Bidder must describe the process that will be adopted to achieve CE marking together with the assurance that will be achieved prior to the start of the Equipment. The Bidder must confirm that CE certification will be provided at an additional cost to the Trust. Bidder must describe how any parts of the Equipment that do not or will not CE mark.</p> <p>Options Provide proof of approval evidence or provide written statement. Provide the proposed evidence that CE marking will be achieved in order for treatment to commence by the start of the Equipment.</p>	0%	0%	0%	Pass/Fail	Fail	
1.1E	<p>Question The Bidder must ensure that the control cabinet complies with the relevant CE marking, or the Bidder must ensure that the control cabinet will achieve CE marking in accordance with the requirements of the relevant CE marking.</p> <p>Response The Bidder must describe the process that will be adopted to achieve CE marking together with the assurance that will be achieved prior to the start of the Equipment. The Bidder must confirm that CE certification will be provided at an additional cost to the Trust. Bidder must describe how any parts of the Equipment that do not or will not CE mark.</p> <p>Options Provide proof of approval evidence or provide written statement. Provide the proposed evidence that CE marking will be achieved in order for treatment to commence by the start of the Equipment.</p>	0%	0%	0%	Pass/Fail	Fail	
1.1F	<p>Question The Bidder must ensure that the control cabinet complies with the relevant CE marking, or the Bidder must ensure that the control cabinet will achieve CE marking in accordance with the requirements of the relevant CE marking.</p> <p>Response The Bidder must describe the process that will be adopted to achieve CE marking together with the assurance that will be achieved prior to the start of the Equipment. The Bidder must confirm that CE certification will be provided at an additional cost to the Trust. Bidder must describe how any parts of the Equipment that do not or will not CE mark.</p> <p>Options Provide proof of approval evidence or provide written statement. Provide the proposed evidence that CE marking will be achieved in order for treatment to commence by the start of the Equipment.</p>	0%	0%	0%	Pass/Fail	Fail	
1.1G	<p>Question The Bidder must ensure that the control cabinet complies with the relevant CE marking, or the Bidder must ensure that the control cabinet will achieve CE marking in accordance with the requirements of the relevant CE marking.</p> <p>Response The Bidder must describe the process that will be adopted to achieve CE marking together with the assurance that will be achieved prior to the start of the Equipment. The Bidder must confirm that CE certification will be provided at an additional cost to the Trust. Bidder must describe how any parts of the Equipment that do not or will not CE mark.</p> <p>Options Provide proof of approval evidence or provide written statement. Provide the proposed evidence that CE marking will be achieved in order for treatment to commence by the start of the Equipment.</p>	0%	0%	0%	Pass/Fail	Fail	
1.1H	<p>Question The Bidder must ensure that the control cabinet complies with the relevant CE marking, or the Bidder must ensure that the control cabinet will achieve CE marking in accordance with the requirements of the relevant CE marking.</p> <p>Response The Bidder must describe the process that will be adopted to achieve CE marking together with the assurance that will be achieved prior to the start of the Equipment. The Bidder must confirm that CE certification will be provided at an additional cost to the Trust. Bidder must describe how any parts of the Equipment that do not or will not CE mark.</p> <p>Options Provide proof of approval evidence or provide written statement. Provide the proposed evidence that CE marking will be achieved in order for treatment to commence by the start of the Equipment.</p>	0%	0%	0%	Pass/Fail	Fail	

- Document 4 - Technical Questionnaire**
- 2.1 Conditions of Supply of Equipment**
- 2.2 Support & Training**
- 2.3 General Facility Environmental Specifications**
- 2.4 Electricity and power & Building Infrastructure**
- 2.5 Equipment Reliability, Efficiency and Uptime Specifications**
- 2.6 Installation & Beam Transport Commissioning**
- 2.7 Radiation Protection**
- 2.8 General Safety / Interlocks**
- 2.9 Control and Monitoring Systems**
- 2.10 Operations, Maintenance and Quality Assurance**
- 2.11 Proton Accelerator and Beamline Parameters**
- 2.12 Clinical Beam Characteristics**
- 2.13 Clinical Cases**
- 2.14 Dosimetry**
- 2.15 Treatment Room Specifications**
- 2.16 In-Room Image Guidance Systems**
- 2.17 Treatment Planning, Verification and Oncology Management Systems**
- 2.18 Upgrades, Innovations and Retrofits**
- 2.19 Fourth Room Option and Research**

- 130 Questions:**
- 31 Pass/Fail
 - 35 For information only (needed for compliant bid)
 - 64 Scored
 - 4 - Excellent confidence
 - 3 - Very good confidence
 - 2 - Good confidence
 - 1 - Minor concerns
 - 0 - Major concerns

Document 4 - Technical Questionnaire

2.1 Conditions of Supply of Equipment

2.2 Support & Training

2.3 General Facility Environmental Specifications

2.4 Electricity and power & Building Infrastructure

2.5 Equipment Reliability, Efficiency and Uptime Specifications

2.6 Installation & Beam Transport Commissioning

2.7 Radiation Protection

2.8 General Safety / Interlocks

2.9 Control and Monitoring Systems

2.10 Operations, Maintenance and Quality Assurance

2.11 Proton Accelerator and Beamline Parameters

2.12 Clinical Beam Characteristics

2.13 Clinical Cases

2.14 Dosimetry

2.15 Treatment Room Specifications

2.16 In-Room Image Guidance Systems

2.17 Treatment Planning, Verification and Oncology Management Systems

2.18 Upgrades, Innovations and Retrofits

2.19 Fourth Room Option and Research

Document 4 - Technical Questionnaire

2.1 Conditions of Supply of Equipment

2.2 Support & Training

2.3 General Facility Environmental Specifications

2.4 Electricity and power & Building Infrastructure

2.5 Equipment Reliability, Efficiency and Uptime Specifications

2.6 Installation & Beam Transport Commissioning

2.7 Radiation Protection

2.8 General Safety / Interlocks

2.9 Control and Monitoring Systems

2.10 Operations, Maintenance and Quality Assurance

2.11 Proton Accelerator and Beamline Parameters

2.12 Clinical Beam Characteristics

2.13 Clinical Cases

2.14 Dosimetry

2.15 Treatment Room Specifications

2.16 In-Room Image Guidance Systems

2.17 Treatment Planning, Verification and Oncology Management Systems

2.18 Upgrades, Innovations and Retrofits

2.19 Fourth Room Option and Research

Document 4 - Technical Questionnaire

2.1 Conditions of Supply of Equipment

2.2 Support & Training

2.3 General Facility Environmental Specifications

2.4 Electricity and power & Building Infrastructure

2.5 Equipment Reliability, Efficiency and Uptime Specifications

2.6 Installation & Beam Transport Commissioning

2.7 Radiation Protection

2.8 General Safety / Interlocks

2.9 Control and Monitoring Systems

2.10 Operations, Maintenance and Quality Assurance

2.11 Proton Accelerator and Beamline Parameters

2.12 Clinical Beam Characteristics

2.13 Clinical Cases

2.14 Dosimetry

2.15 Treatment Room Specifications

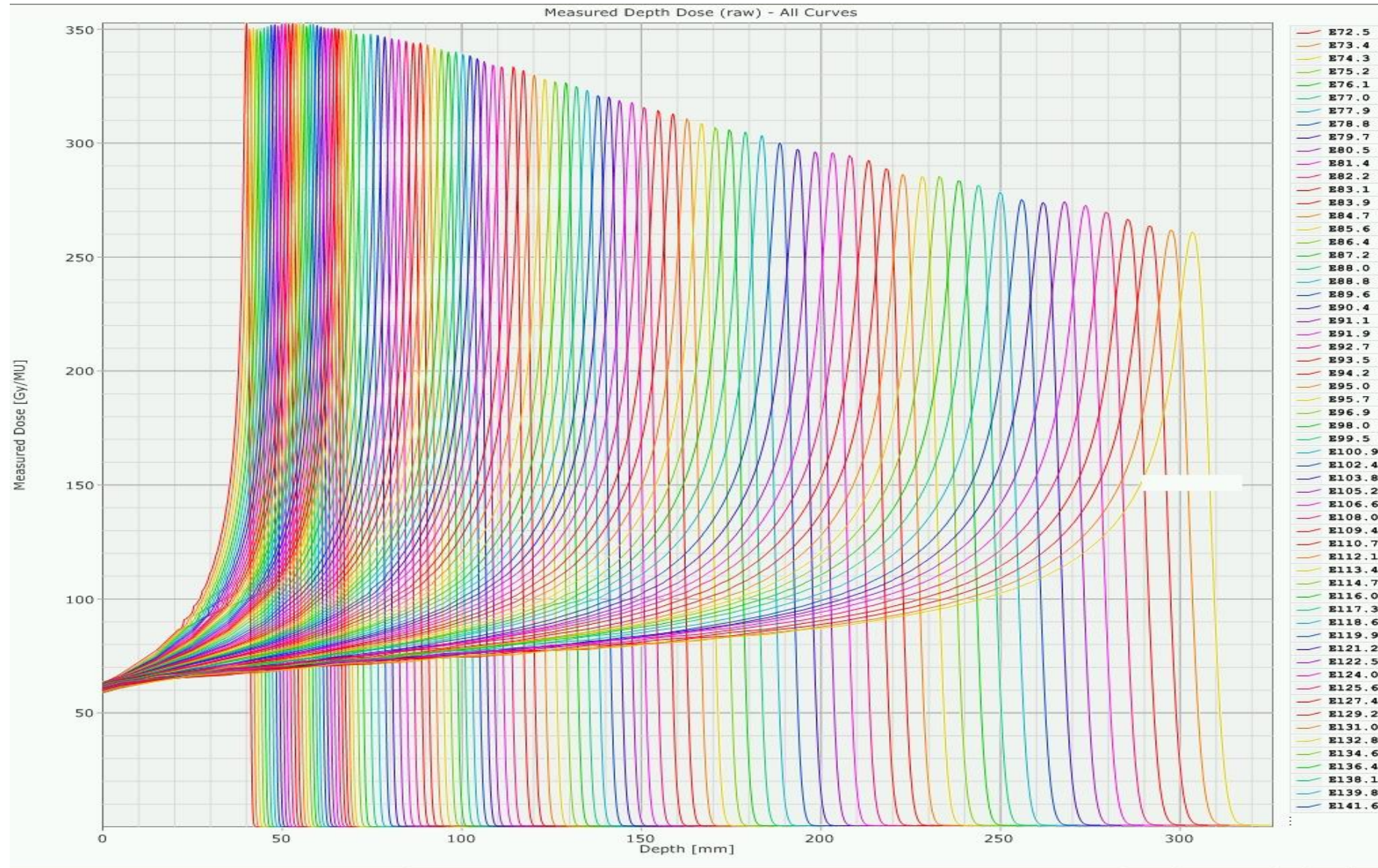
2.16 In-Room Image Guidance Systems

2.17 Treatment Planning, Verification and Oncology Management Systems

2.18 Upgrades, Innovations and Retrofits

2.19 Fourth Room Option and Research

Monte Carlo generated pre-commissioning data at MD Anderson



Document 4 - Technical Questionnaire

2.1 Conditions of Supply of Equipment

2.2 Support & Training

2.3 General Facility Environmental Specifications

2.4 Electricity and power & Building Infrastructure

2.5 Equipment Reliability, Efficiency and Uptime Specifications

2.6 Installation & Beam Transport Commissioning

2.7 Radiation Protection

2.8 General Safety / Interlocks

2.9 Control and Monitoring Systems

2.10 Operations, Maintenance and Quality Assurance

2.11 Proton Accelerator and Beamline Parameters

2.12 Clinical Beam Characteristics

2.13 Clinical Cases

2.14 Dosimetry

2.15 Treatment Room Specifications

2.16 In-Room Image Guidance Systems

2.17 Treatment Planning, Verification and Oncology Management Systems

2.18 Upgrades, Innovations and Retrofits

2.19 Fourth Room Option and Research

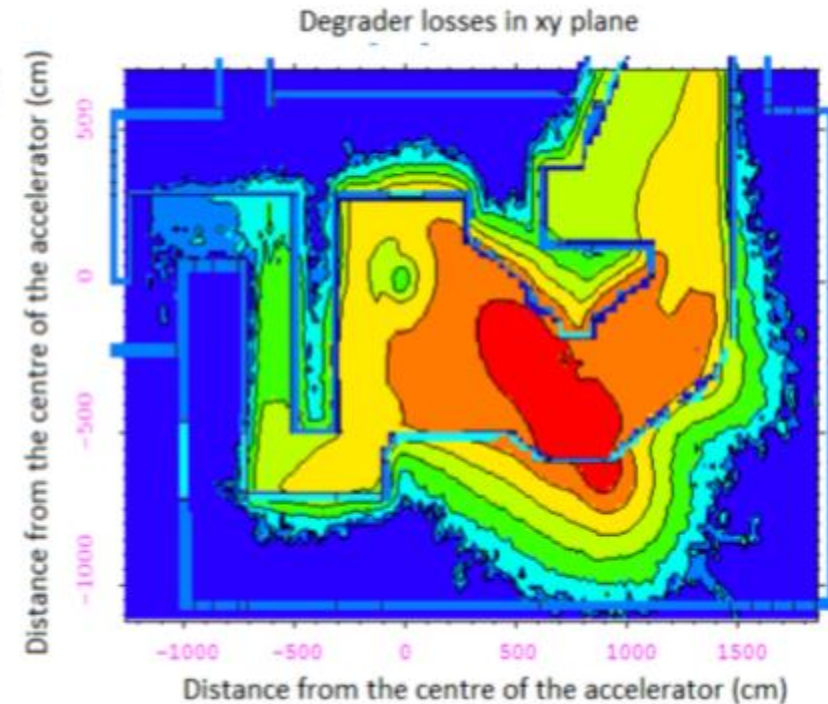
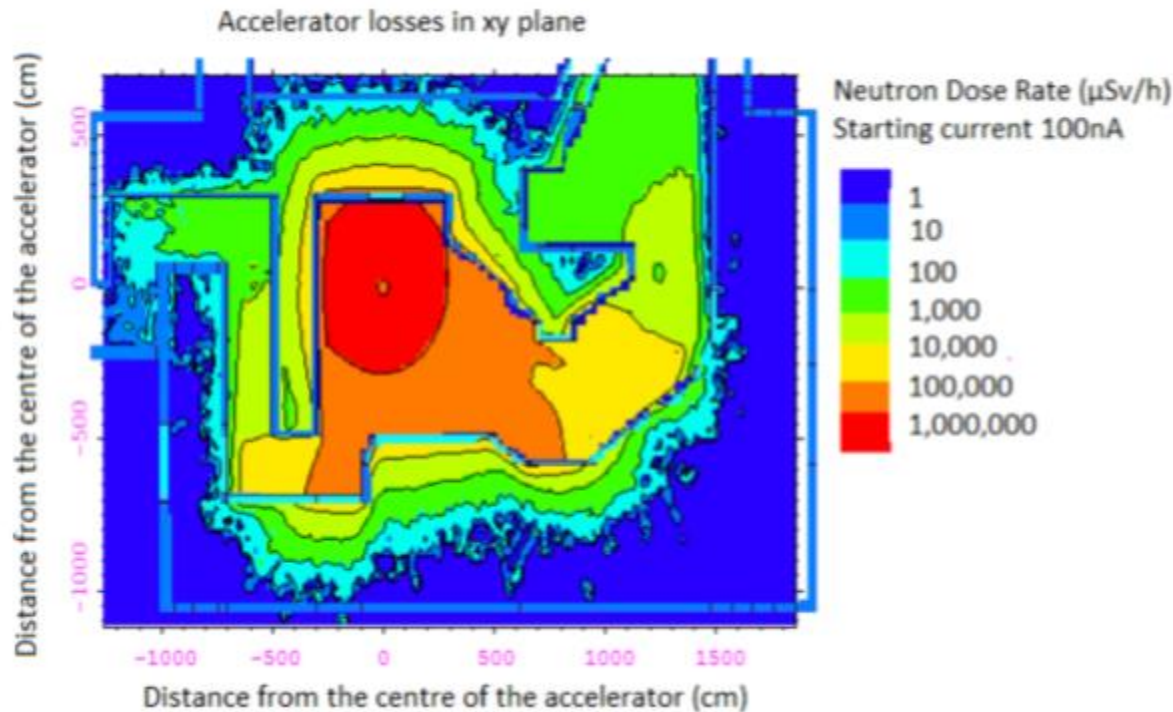
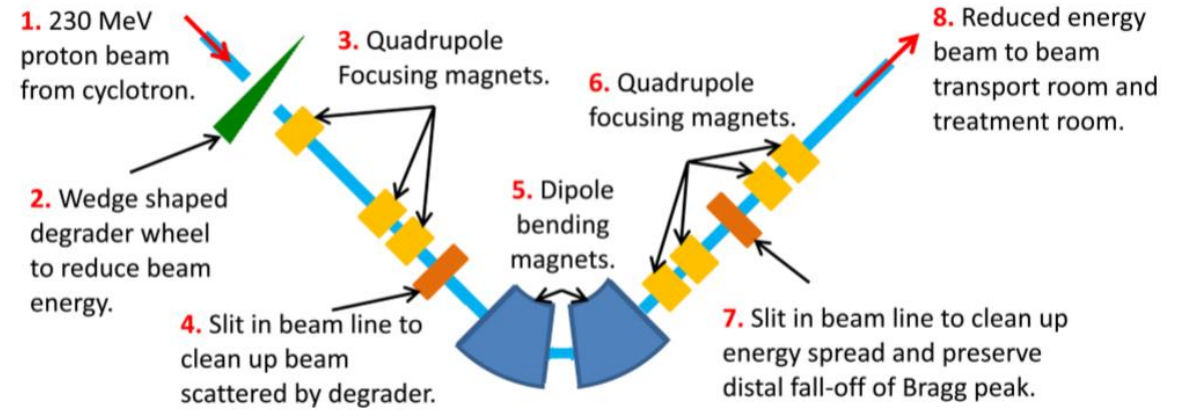
Workload calculations based on estimates of:

- Patient throughput
- Numbers and angles of beams
- Range of energies used
- Volume of targets

Activation of components, incl. air and water.

Ducting in shielding walls – BIM.

Monte Carlo modelling of losses....



*Particle
Therapy
Co-
Operative
Group*

PTCOG Report 1

PTCOG Publications Sub-Committee Task Group on Shielding Design and Radiation Safety of Charged Particle Therapy Facilities

Series in Physics and Engineering in Medicine and Biology

SECOND
EDITION

Design and Shielding of Radiotherapy Treatment Facilities

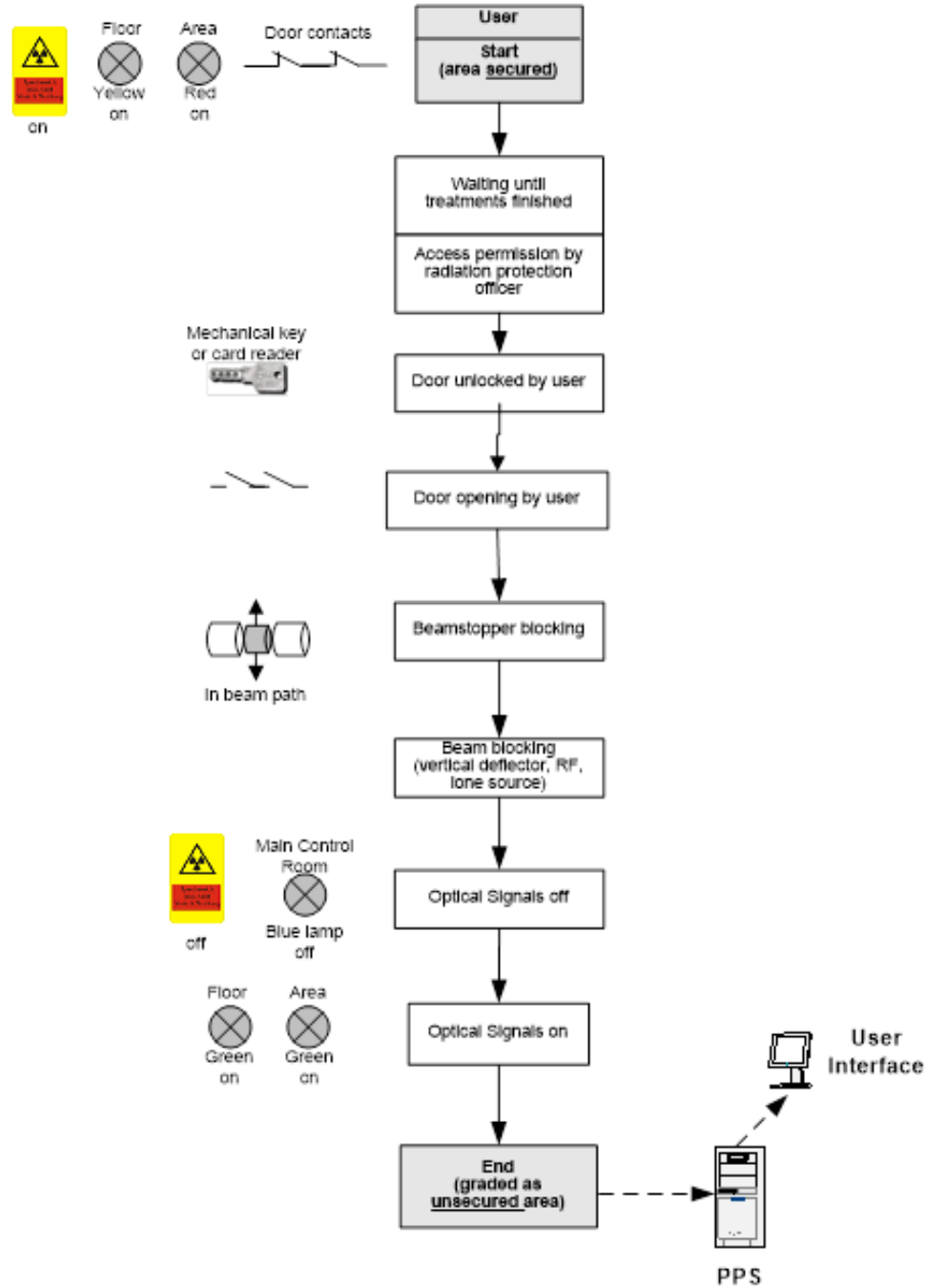
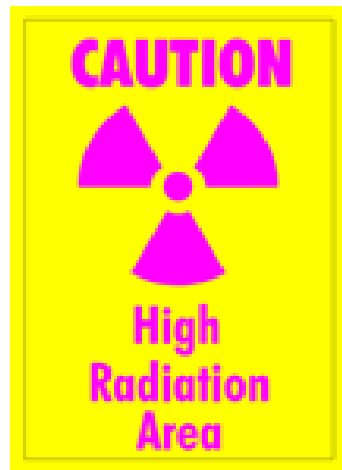
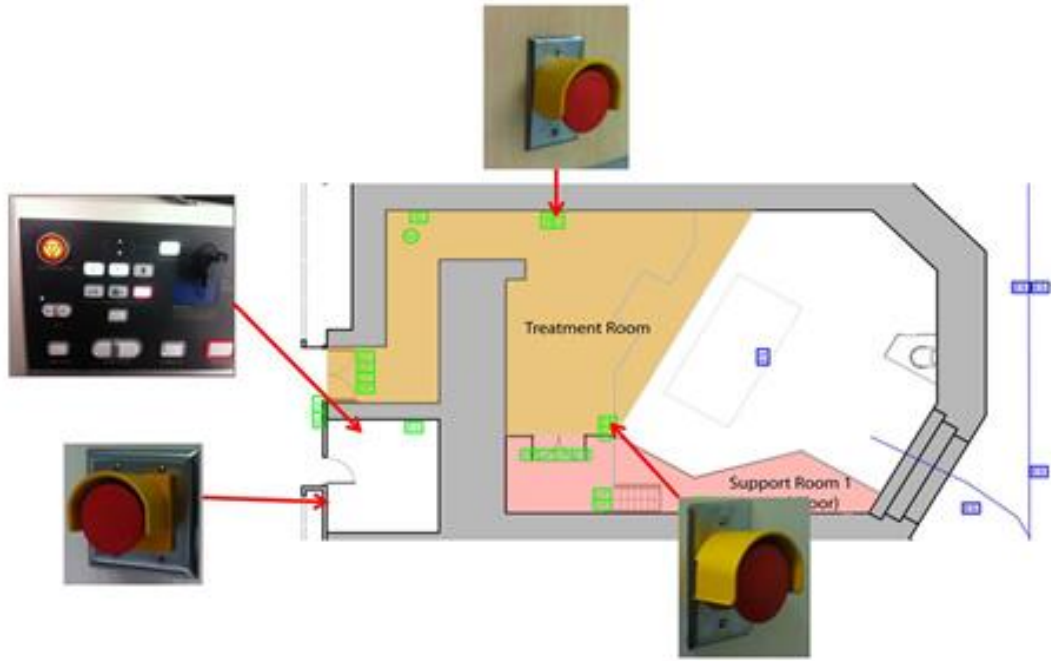
IPEM Report 75

Chapter 11

Radiation shielding and safety for particle
therapy facilities

R L Maughan, M J Hardy, M J Taylor, J Reay and R Amos

Radiation and general safety interlock system



Document 4 - Technical Questionnaire

2.1 Conditions of Supply of Equipment

2.2 Support & Training

2.3 General Facility Environmental Specifications

2.4 Electricity and power & Building Infrastructure

2.5 Equipment Reliability, Efficiency and Uptime Specifications

2.6 Installation & Beam Transport Commissioning

2.7 Radiation Protection

2.8 General Safety / Interlocks

 **2.9 Control and Monitoring Systems**

2.10 Operations, Maintenance and Quality Assurance

2.11 Proton Accelerator and Beamline Parameters

2.12 Clinical Beam Characteristics

2.13 Clinical Cases

 **2.14 Dosimetry**

2.15 Treatment Room Specifications

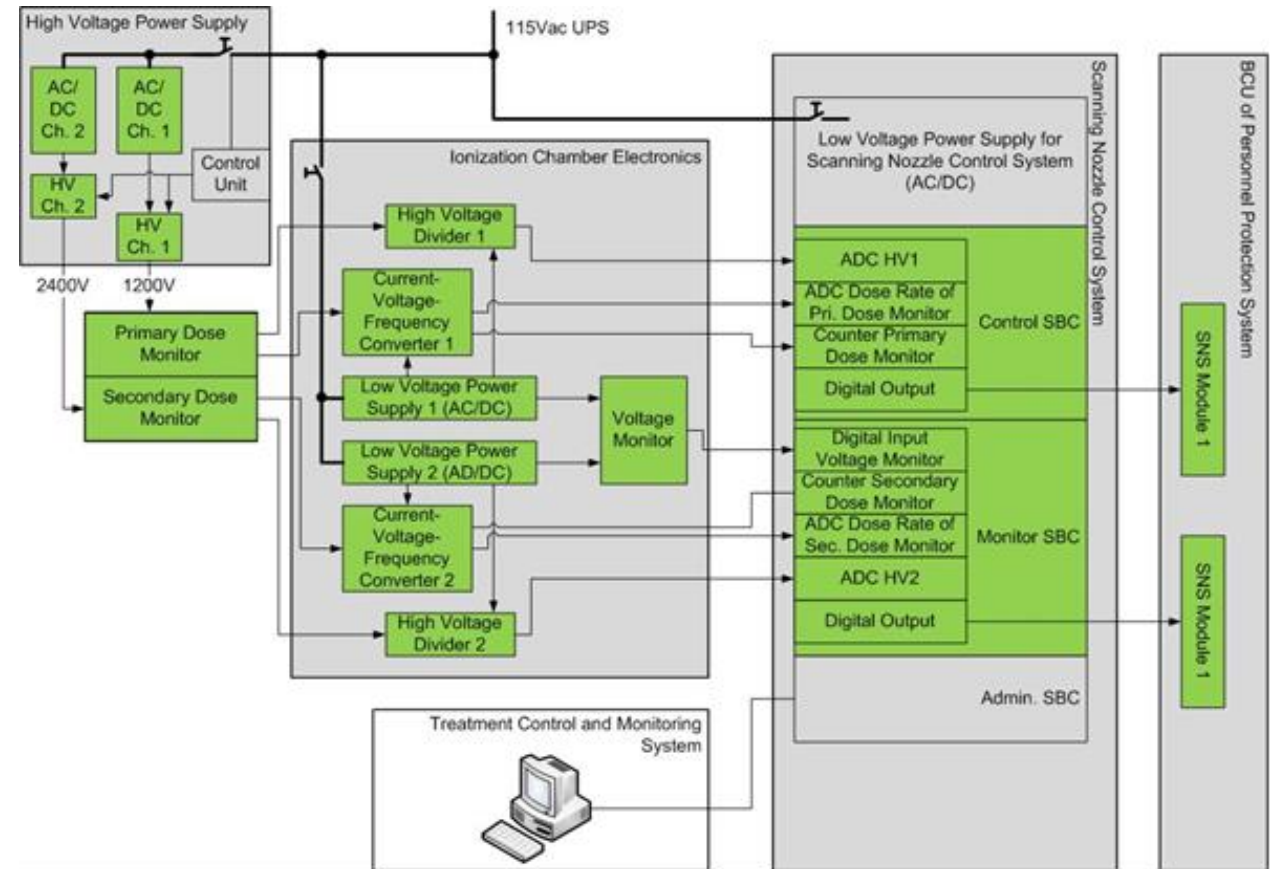
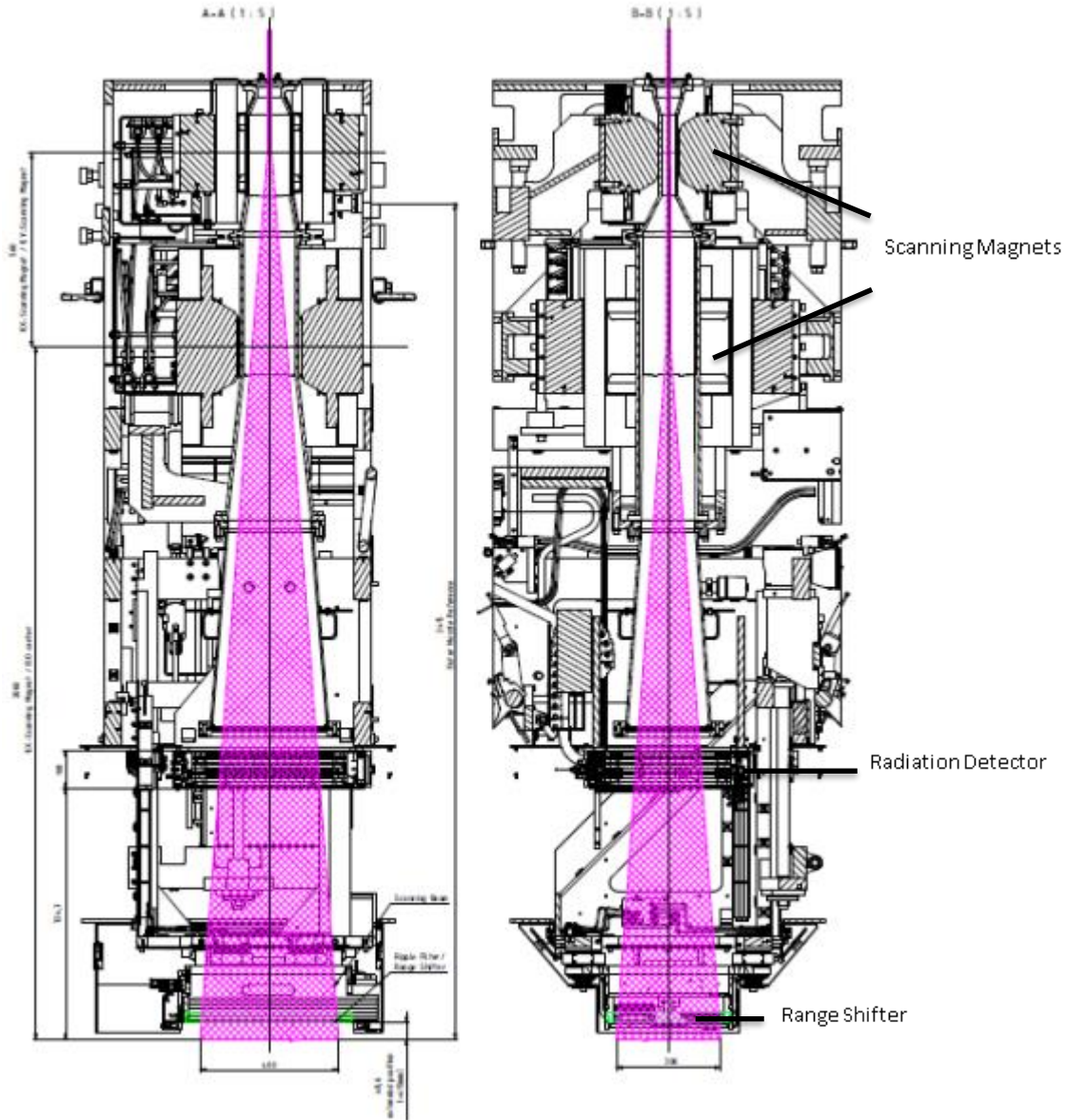
2.16 In-Room Image Guidance Systems

2.17 Treatment Planning, Verification and Oncology Management Systems

2.18 Upgrades, Innovations and Retrofits

2.19 Fourth Room Option and Research

Dosimetry, telemetry, and beam control systems



Document 4 - Technical Questionnaire

2.1 Conditions of Supply of Equipment

2.2 Support & Training

2.3 General Facility Environmental Specifications

2.4 Electricity and power & Building Infrastructure

2.5 Equipment Reliability, Efficiency and Uptime Specifications

2.6 Installation & Beam Transport Commissioning

2.7 Radiation Protection

2.8 General Safety / Interlocks

2.9 Control and Monitoring Systems

2.10 Operations, Maintenance and Quality Assurance

2.11 Proton Accelerator and Beamline Parameters

 **2.12 Clinical Beam Characteristics**

2.13 Clinical Cases

2.14 Dosimetry

2.15 Treatment Room Specifications

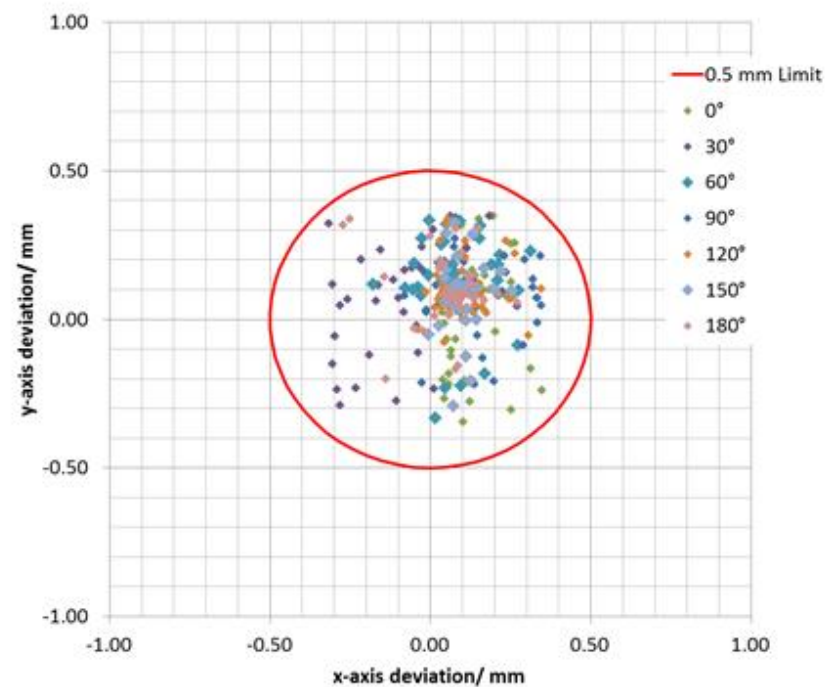
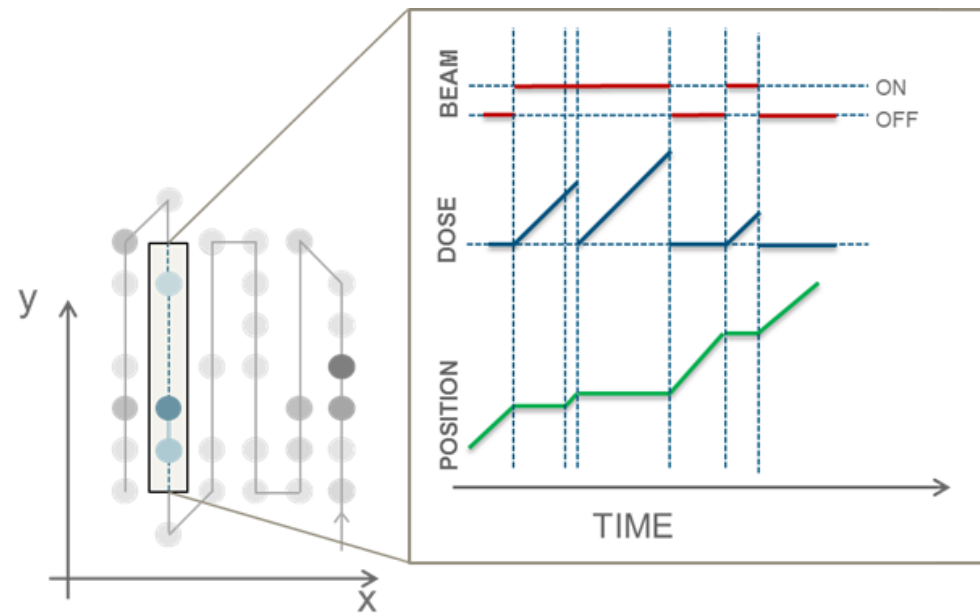
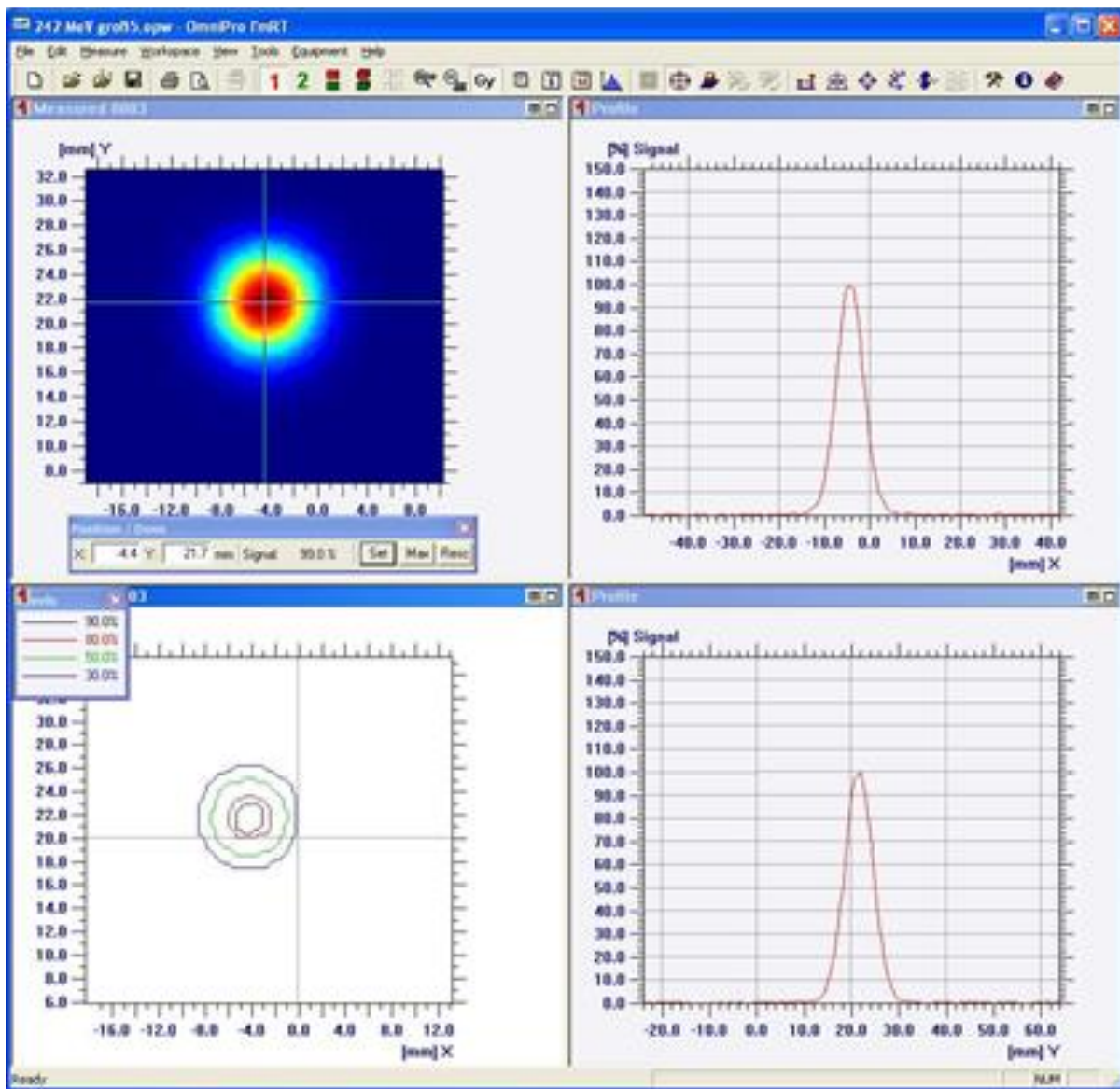
2.16 In-Room Image Guidance Systems

2.17 Treatment Planning, Verification and Oncology Management Systems

2.18 Upgrades, Innovations and Retrofits

2.19 Fourth Room Option and Research

Clinical beam characteristics



Document 4 - Technical Questionnaire

2.1 Conditions of Supply of Equipment

2.2 Support & Training

2.3 General Facility Environmental Specifications

2.4 Electricity and power & Building Infrastructure

2.5 Equipment Reliability, Efficiency and Uptime Specifications

2.6 Installation & Beam Transport Commissioning

2.7 Radiation Protection

2.8 General Safety / Interlocks

2.9 Control and Monitoring Systems

2.10 Operations, Maintenance and Quality Assurance

2.11 Proton Accelerator and Beamline Parameters

2.12 Clinical Beam Characteristics

2.13 Clinical Cases

2.14 Dosimetry

2.15 Treatment Room Specifications

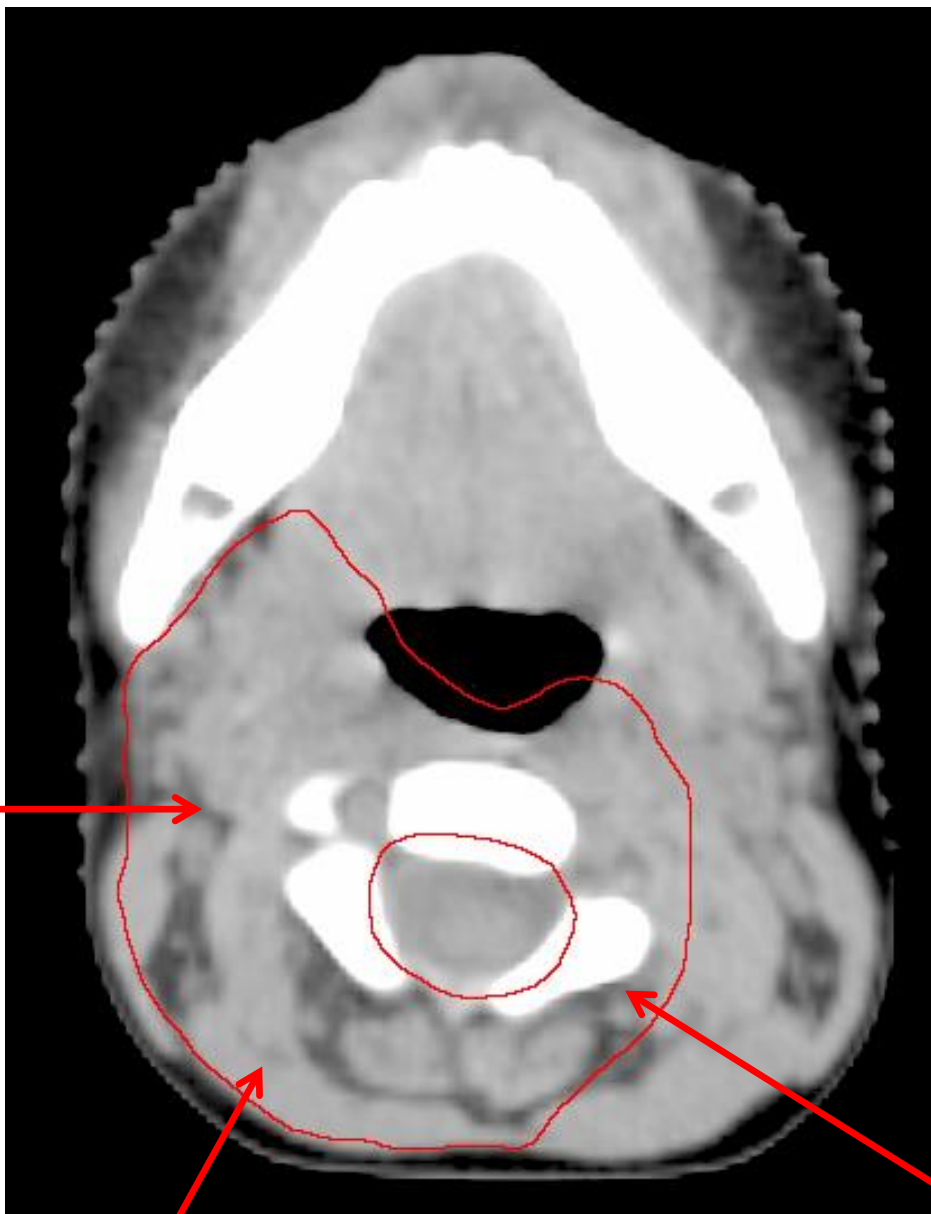
2.16 In-Room Image Guidance Systems

2.17 Treatment Planning, Verification and Oncology Management Systems

2.18 Upgrades, Innovations and Retrofits

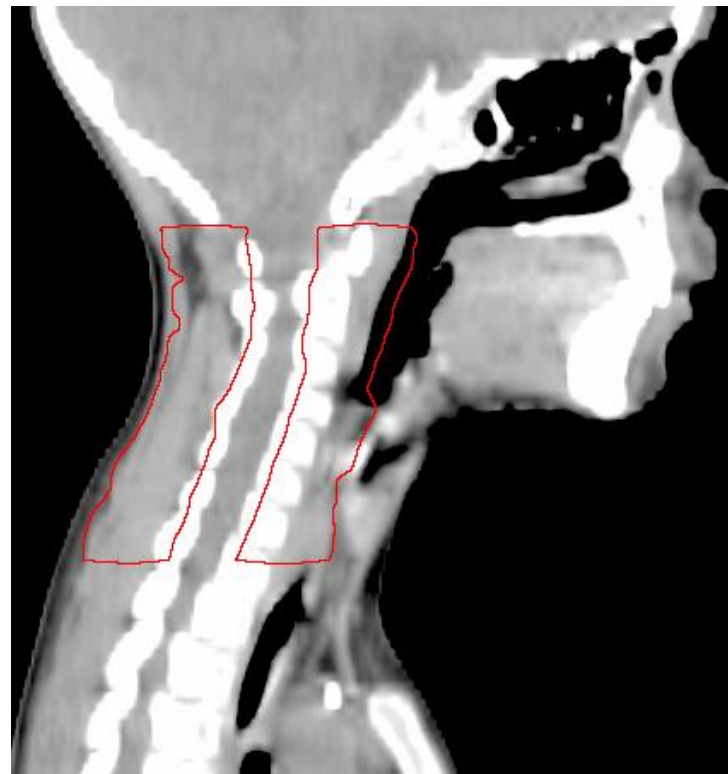
2.19 Fourth Room Option and Research

Clinical Case # 1



Ewing Sarcoma Right C-Spine

- Rx: 54 Gy (RBE) / 30 fx to C-spine and right neck (*PTV ED*)
- IMPT with component of SFUD (*at least 20% IMPT*)
- RPO, gantry angle $\sim 200 - 210$ deg
- LPO, gantry angle $\sim 130 - 140$ deg
- Rt Lat, gantry angle 90 deg
- Same angles for SFUD and IMPT components

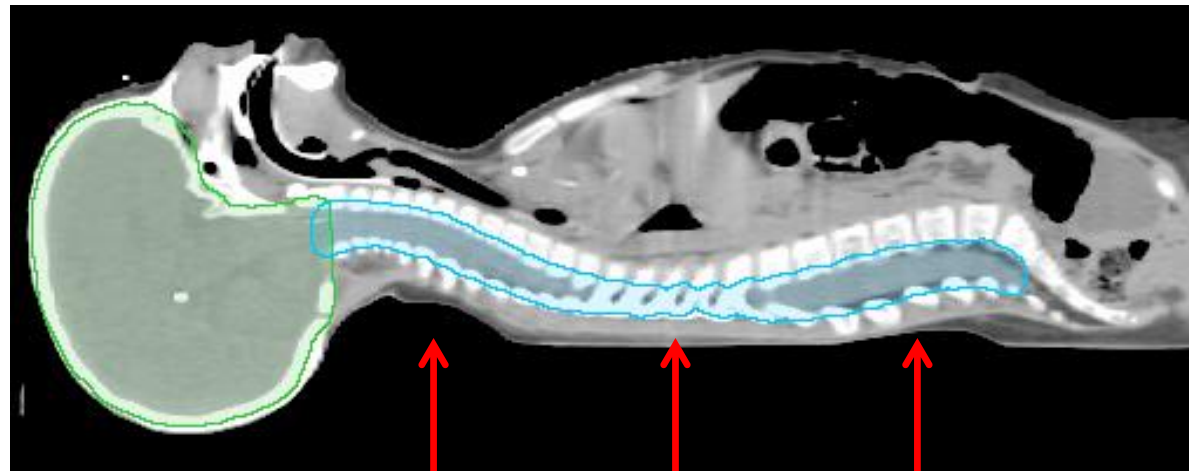
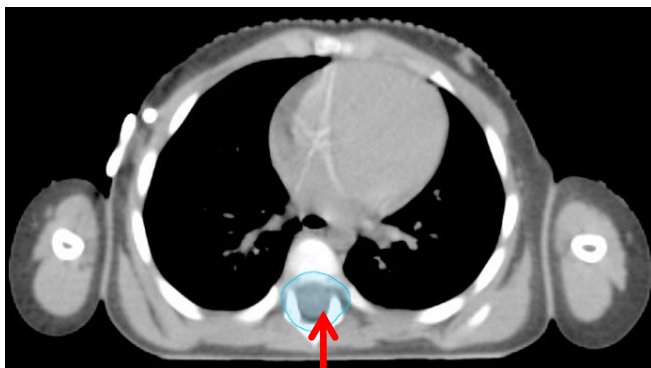
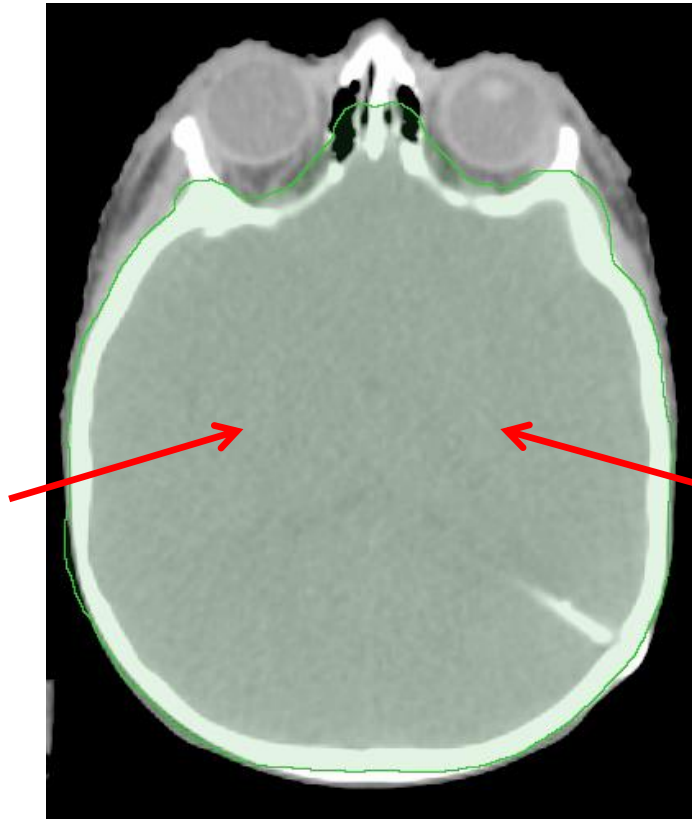


Clinical Case # 2 (Phase 1)

Metastatic Ependymoma

- IMPT with component of SFUD (*at least 20% IMPT*)
- Patient under GA in supine position

- Phase 1 Rx: 36 Gy (RBE) / 20 fx to whole CNS (*brain + spine PTV*)
- Multiple isocenters**
- RPO and LPO fields to the brain, gantry angles ~ 255 and 105 deg respectively
- Multiple PA fields to spine (*number of fields depends on field size limitations*)
- IMPT component to taper field matches to improve plan robustness

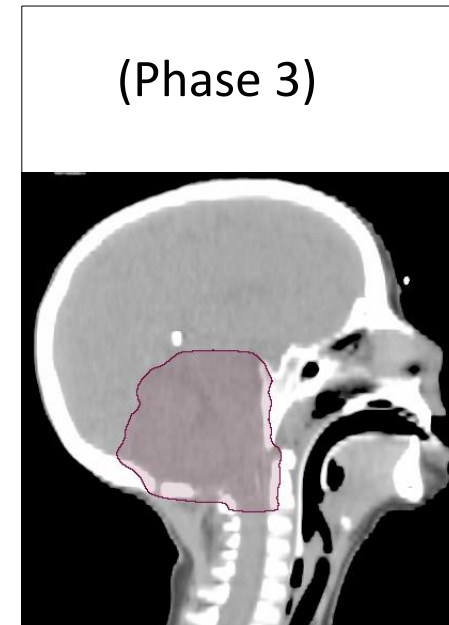
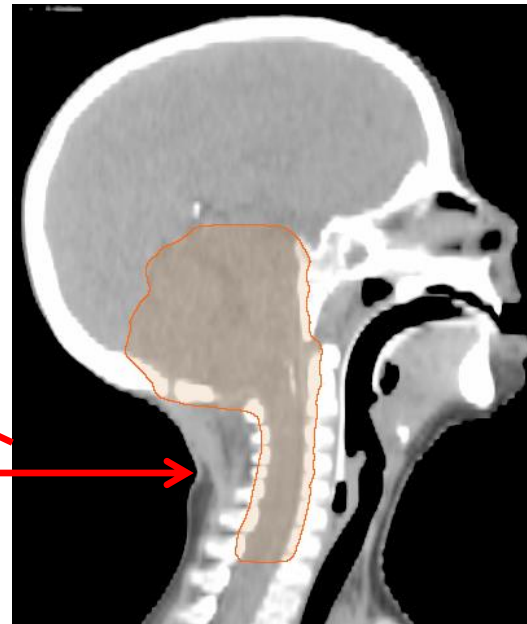


Clinical Case # 2 (Phases 2 & 3)

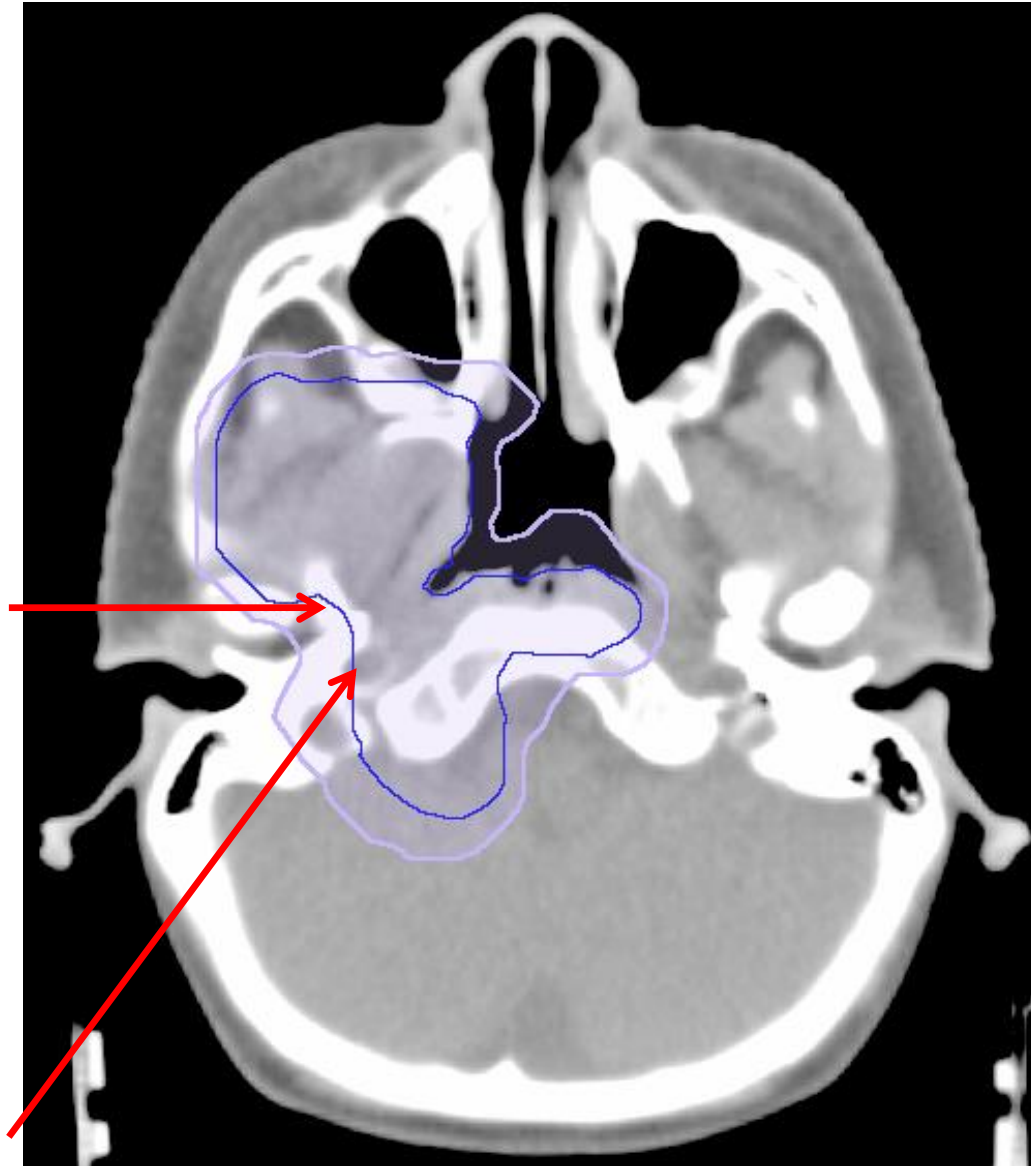


Metastatic Ependymoma

- IMPT with component of SFUD (*at least 20% IMPT*)
- Patient under **GA** in **supine** position
- Phase 2 Rx: 9 Gy (RBE) / 5 fx to C-spine and post fossa (*PTVPh2*)
- LPO, gantry angle ~ 120 – 130 deg
- RPO, gantry angle ~ 230 – 240 deg
- PA, gantry angle = 180 deg
- IMPT – eg, contribution of PA only at C-spine level
- Phase 3 Rx: 9 Gy (RBE) / 5 fx to post fossa (*PTVPh3*)
- Repeat LPO and RPO IMPT fields
- Omit PA component

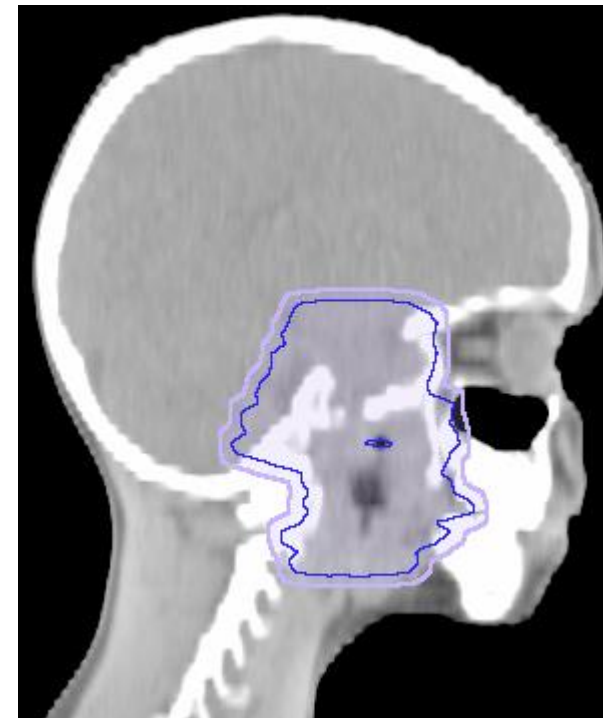


Clinical Case # 3



Alveolar Rhabdomyosarcoma (Post nasal space)

- Rx: 50.4 Gy (RBE) / 28 fx to PTV
- IMPT with component of SFUD (*at least 20% IMPT*)
- RPO, gantry angle ~ 220 deg
- Rt Lat, (*or very slight RPO*)
- SFUD up to a dose of approx. 40 Gy (RBE), then IMPT to better spare the OARs. Same gantry angles for SFUD and IMPT



Clinical Case # 1

CRITICAL ORGAN LABEL	DOSE CONSTRAINT [Gy (RBE)]
Canal (PRV)	50 Gy (RBE)
Parotid (volume NOT PRV)	24 Gy (RBE) max; 18 Gy (RBE) mean
PharynxConstrictors (Volume NOT PRV)	50 Gy (RBE) mean but do not compromise PTV dose
Mandible	Either: < 10 Gy (RBE) whole structure OR: > 10 Gy (RBE) but < 30 Gy (RBE) uniformly over whole structure.

Clinical Case # 2

CRITICAL ORGAN LABEL	DOSE CONSTRAINT [Gy (RBE)]
Pituitary	For record only
Eyes	54 Gy (RBE)
Chiasm	55 Gy (RBE)
Brainstem	55 Gy (RBE)
Optic Nerves	45 Gy (RBE)
Lens	5 Gy (RBE)
Parotid	Mean dose 24 Gy (RBE) to organ
Facial bones	Either: < 10 Gy (RBE) whole structure OR: > 10 Gy (RBE) but < 30 Gy (RBE) uniformly over whole structure.

Clinical Case # 3

CRITICAL ORGAN LABEL	DOSE CONSTRAINT [Gy (RBE)]
RIGHT EYE	Maximum Dose <45 Gy (RBE) to organ
RIGHT LENS	Maximum Dose <5 Gy (RBE) to organ
RIGHT OPTIC NERVE	Maximum Dose <50 Gy (RBE) to organ PRV
LEFT EYE	Maximum Dose <45 Gy (RBE) to organ
LEFT LENS	Maximum Dose <5 Gy (RBE) to organ
LEFT OPTIC NERVE	Maximum Dose <50 Gy (RBE) to organ PRV
OPTIC CHIASM	Maximum Dose <55 Gy (RBE) to organ PRV
BRAINSTEM	Maximum Dose <55 Gy (RBE) to organ PRV
LEFT PAROTID	Maximum Mean dose 24 Gy (RBE) to organ
Mandible	Either: < 10 Gy (RBE) whole structure OR: > 10 Gy (RBE) but < 30 Gy (RBE) uniformly over whole structure.

Plans returned as DICOM files and evaluated for clinical acceptability

Document 4 - Technical Questionnaire

2.1 Conditions of Supply of Equipment

2.2 Support & Training

2.3 General Facility Environmental Specifications

2.4 Electricity and power & Building Infrastructure

2.5 Equipment Reliability, Efficiency and Uptime Specifications

2.6 Installation & Beam Transport Commissioning

2.7 Radiation Protection

2.8 General Safety / Interlocks

2.9 Control and Monitoring Systems

2.10 Operations, Maintenance and Quality Assurance

2.11 Proton Accelerator and Beamline Parameters

2.12 Clinical Beam Characteristics

2.13 Clinical Cases

2.14 Dosimetry

2.15 Treatment Room Specifications

2.16 In-Room Image Guidance Systems

2.17 Treatment Planning, Verification and Oncology Management Systems

2.18 Upgrades, Innovations and Retrofits

2.19 Fourth Room Option and Research



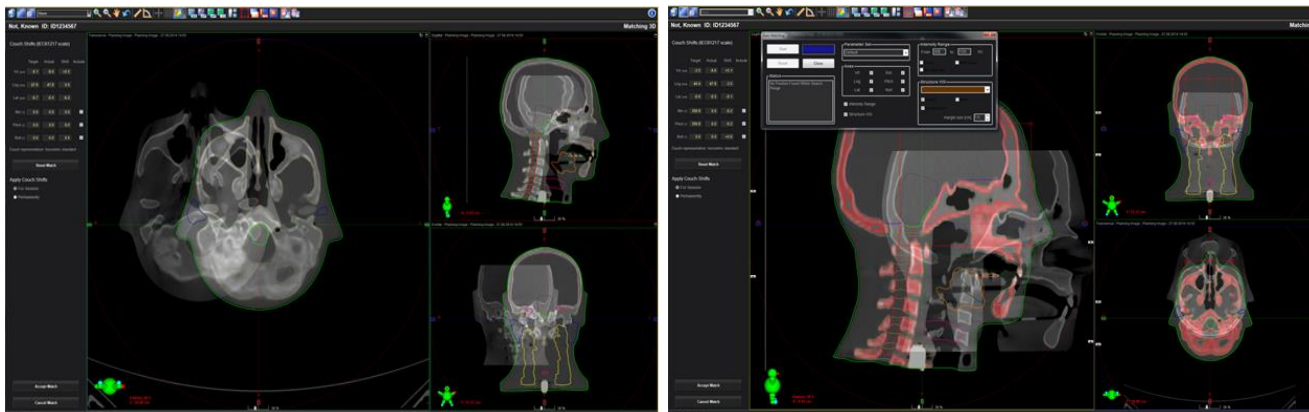


Treatment room specs:

- Patient positioner specs
 - Compatibility with immobilization
- In-room telemetry/audio
- Collision detection system
- Anesthetic gases
- Functionality/workflow
-

Image-guidance:

- Laser alignment systems
- 2D/3D/4D imaging
- Acquisition and reconstruction speeds
- Image quality
- Connectivity to auxiliary systems
 - Optical surface imaging
 - Tracking



Accommodations

- Patient pathway through facility
- Waiting area
- Consultation rooms
- Anaesthesia – recovery
- Imaging suite – CT (DECT), MRI, PET/CT,...
- Staff accommodations
- Teleconference rooms
- Storage – e.g. physics QA equipment
-

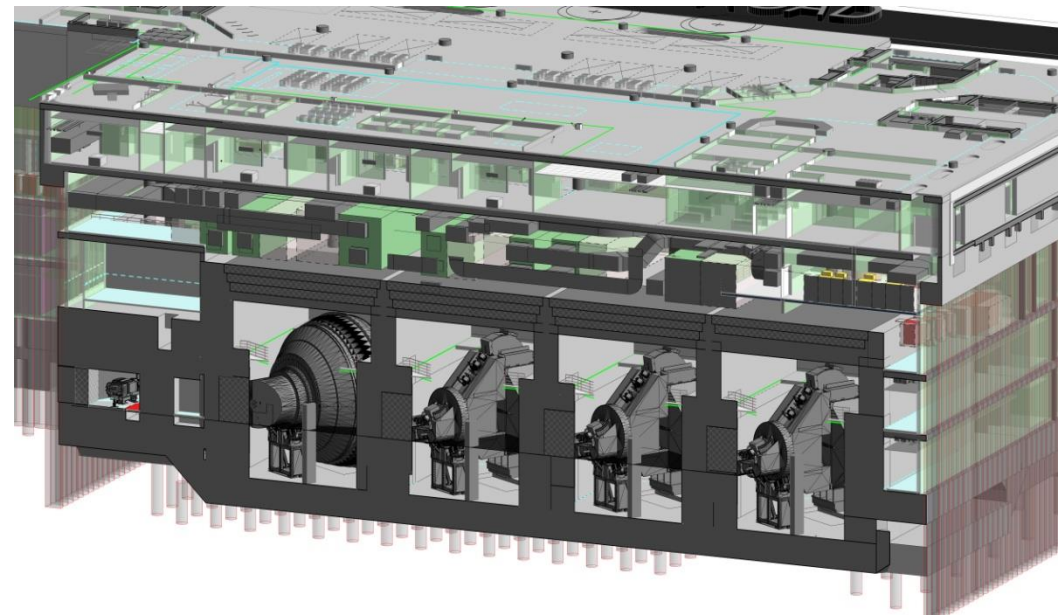
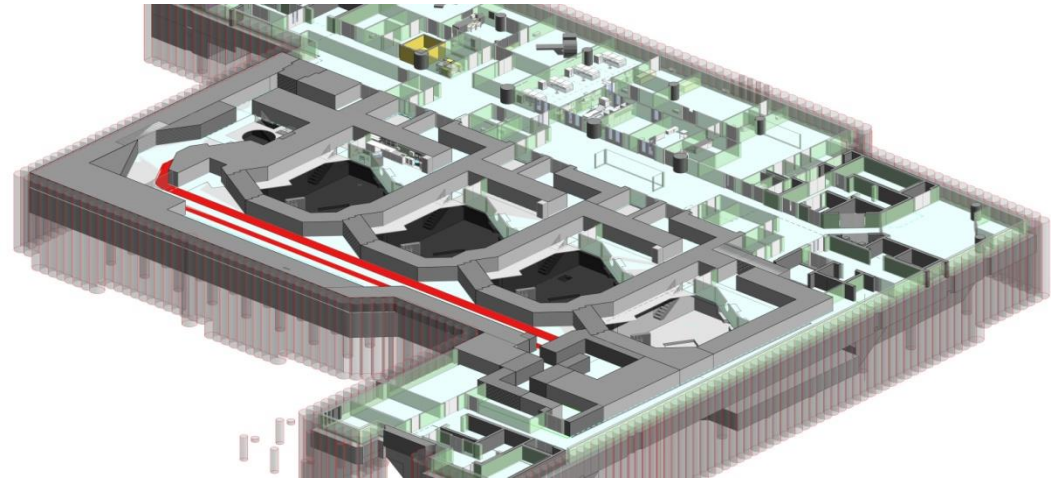
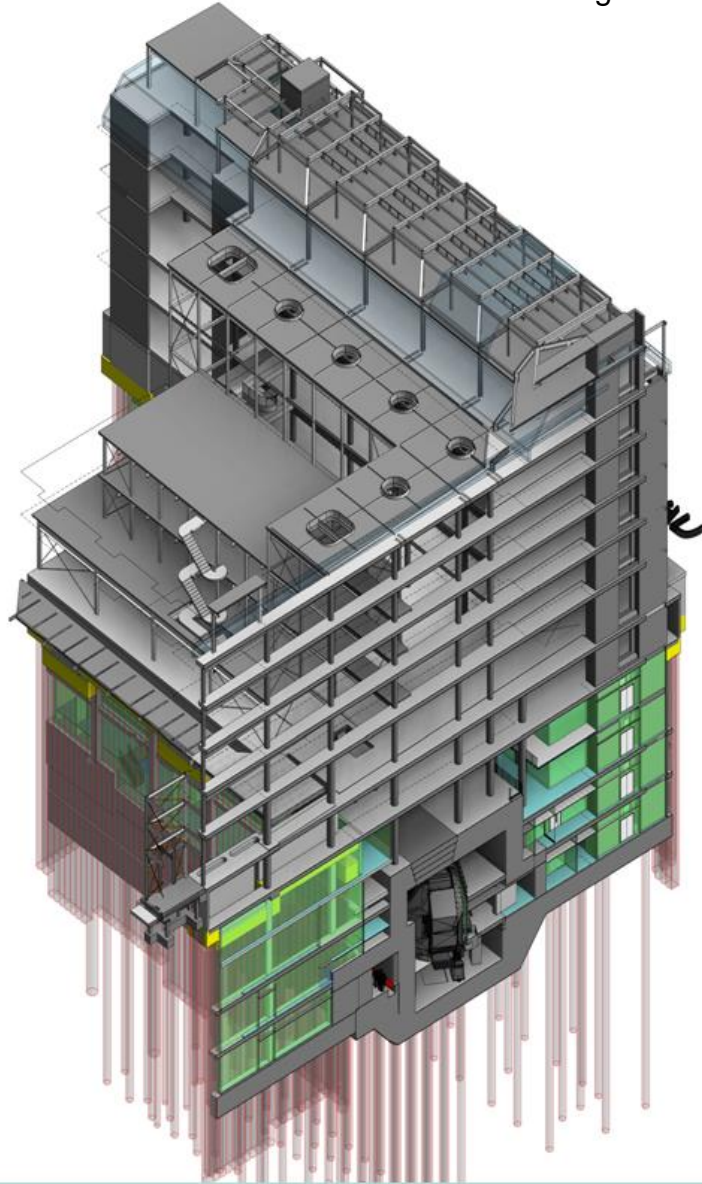
Other infrastructure

- Workforce – numbers and training
- Education of referrers and public
- Referral pathways
- Outcomes data collection and storage
- Academic and industrial partnerships
-

Have people involved who have experience and understand of clinical radiotherapy

Zakrzewska P, Pitt M, Amos RA, D'Souza D & Ahmed T.
Application of building information modelling (BIM) in the design, construction, and operations management of a complex proton beam therapy facility in central London.

Proceedings of PTCOG 54. *Int J Particle Ther.* 2015;2(1):331-332



Thank you!

