

ALGORITHMS- QA & COMMISSIONING



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Disclosure: None



AAPM SPRING CLINICAL MEETING, MARCH 30 - APRIL 2, 2019

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EDUCATIONAL OBJECTIVES

To review the minimum requirements for TPS dose algorithm commissioning & QA.

To review the important issues for consideration during data acquisition, beam modelling, and validation tests.



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MANY KEY DOCUMENTS AND RESOURCES

Just list a few:

- TG53: TPS acceptance, commission and on-going QA.
- TG106: beam data commissioning equipment and procedures.
- IAEA technical report #430: commissioning and QA of TPS.
- TG65: tissue inhomogeneity corrections
- TG119: guidance document on IMRT
- MPPG5A: commissioning and QA of treatment planning dose calculations
- And many many more...



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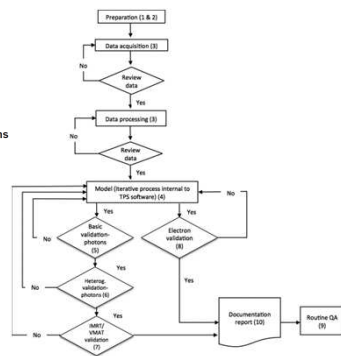
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AAPM MEDICAL PHYSICS PRACTICE GUIDELINE 5.A.: COMMISSIONING AND QA OF TREATMENT PLANNING DOSE CALCULATIONS

JOURNAL OF APPLIED CLINICAL MEDICAL PHYSICS, VOLUME 16, NUMBER 5, 2015

AAPM Medical Physics Practice Guideline 5.a.: Commissioning and QA of Treatment Planning Dose Calculations — Megavoltage Photon and Electron Beams

Medical Physics Practice Guideline. Jennifer B. Smolowitz, Chair,
Indira J. Das, Vladimir Fejgelman, Benedict A. Fraass, Stephen F. Kry,
Ingrid R. Marshall, Dimitris N. Mihailidis, Zoubi Ouhbi, Timothy Ritter,
Michael G. Snyder, Lynne Farrow, AAPM Staff



TIME REQUIRED FOR COMMISSIONING

Assuming 12–16 QMP work hours per day (1.5 to 2.0 FTEs), reasonable time estimates are

- two to four weeks for a single energy photon beam
- six to eight weeks for two photon energies and five electron energies.
- Addition of a second algorithm for a given beam will increase commissioning time and effort.

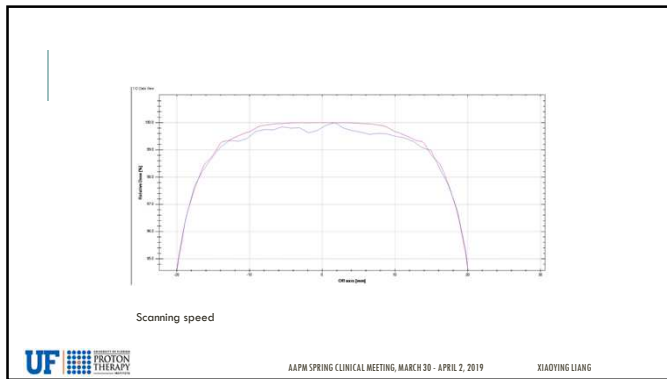
This will depend strongly on how much commissioning data need to be collected and the availability and experience of the QMP(s) involved, the adequacy and availability of the equipment used, and the access to the accelerator

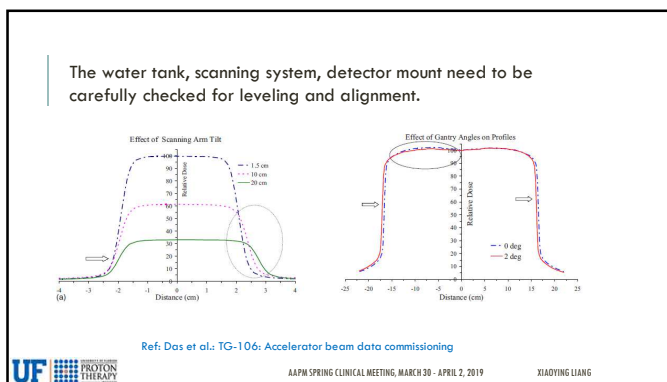
Ref: MPPG5A



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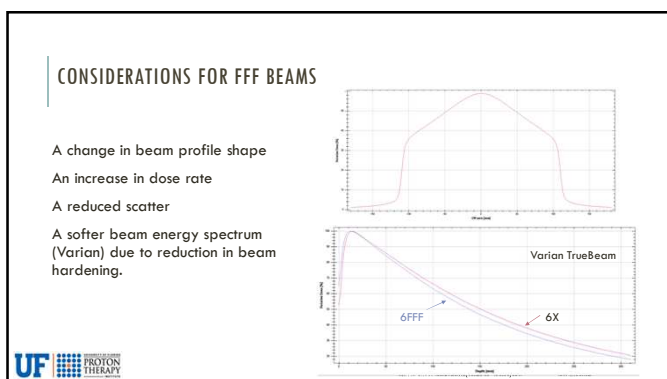


Table 1. Beam characteristics for 6 and 10 MV FFF and cFF beams from Varian and Elekta. See Xiao et al (2015) for Siemens data.

		Nominal energy (MV)	Filtration	Effective energy (MV) ^a	d_{max} (cm)	D_{10} (%)	TPR ₂₀₀	Max dose rate (MU min ⁻¹)	Dose (mGy) per pulse ^b
Varian	FFF	6	0.8 mm	4	1.3	64.2	0.630	1400	0.8
		10	Brass plate	8	2.2	71.7	0.705	2400	1.3
	cFF	6	6/10 MV flattening filter	6	1.4	66.4	0.666	600	0.3
Elekta	FFF	6	2.0 mm stainless steel plate	6	1.7	67.5	0.684	1400	0.6
		10		10	2.4	73.0	0.734	2200	0.9
	cFF	6	6/10 MV flattening filter	6	1.5	67.5	0.678	600	0.2
		10		10	2.1	73.0	0.721	600	0.4

^aClinical effective energy, based on TPR₂₀₀ and percentage depth dose falloff.

^bMeasured at d_{max} on beam central axis for standard reference conditions.

Note: D_{max} refers to depth of maximum dose, MU are monitor units.

Ref: G Budgell et al. IPeM topical report 1: guidance on implementing flattening filter free (FFF) radiotherapy. PMB. 61 (2016) 8360



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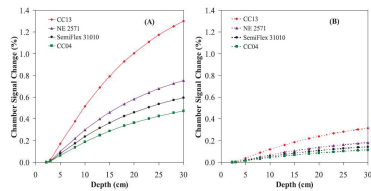


Figure 4. Effect on ionisation chamber signal at depth due to recombination losses for various detectors for (A) 10 MV FFF photons and (B) 10 MV flattened photons for a Varian TrueBeam. Measurements are for a 10 × 10 cm field at 100 cm SSD, using an operating potential of ~350V for all chambers. The ion chamber volumes are respectively: IBA CC13 0.13 cm³, NE 2571 0.6 cm³, PTW 31010 0.125 cm³, IBA CC34 0.04 cm³.

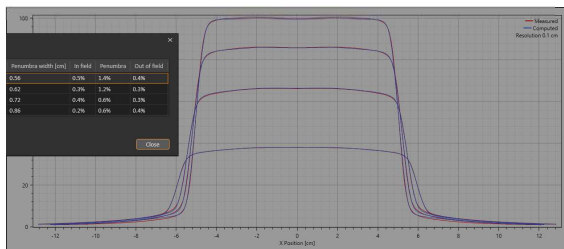
G Budgell et al. Phys. Med. Biol. 61 (2016) 8360



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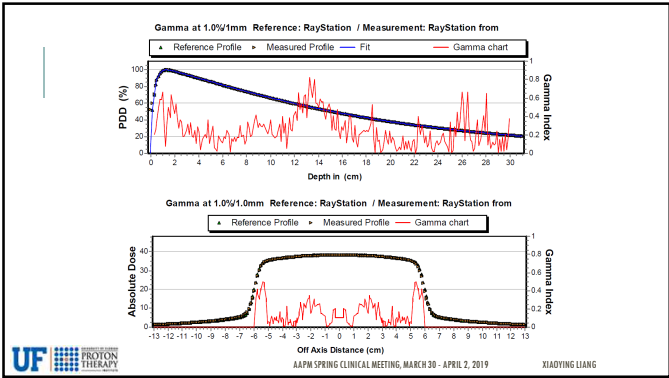
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MODELLING



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VALIDATION TESTS

- ❑ Non measurement "sanity checks"
- ❑ Basic photon beam validation
- ❑ Heterogeneity correction validation
- ❑ IMRT/VMAT
- ❑ E2E test

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VALIDATION TESTS I: NON-MEASUREMENT "SANITY" CHECK

Field configurations are the same as those used for modeling

TABLE 3. TPS model comparison tests and tolerances. **MPPG5A**

Test	Comparison	Description	Tolerance
5.1	Dose distributions in planning module vs. modeling (physics) module	Comparison of dose distribution for large (> 30x30cm ²) field.	Identical ^a
5.2	Dose in test plan vs. clinical calibration condition ^b	Reference calibration condition check	0.5%
5.3	Dose distribution calculated in planning system vs. commissioning data	PDD and off axis output factors for a large and a small field size	2%

^a Identical to within the expected statistical uncertainty (considering noise and calculation grid size).
^b TPS absolute dose at reference point.

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VALIDATION TESTS II –BASIC PHOTON BEAM VALIDATION

Validation tests for

- clinical relevant SSD.
- field shaping using the MLC with jaws at clinical relevant position.
- oblique beam angles
- Wedges

measurements in the high-dose region, penumbra, and low-dose tail regions should be compared to calculated values at various depths.

TABLE 5. Basic TPS photon beam evaluation methods and tolerances. MPPG5A

Region	Evaluation Method	Tolerance ^a (consistent with IROC Houston)
High dose	Relative dose with one parameter change from reference conditions	2%
	Relative dose with multiple parameter changes ^b	5%
Penumbra	Distance to agreement	3 mm
Low-dose tail	Up to 5 cm from field edge	3% of maximum field dose

³ Tolerances are relative to local dose unless otherwise noted

^b For example, off-axis with physical wedge.

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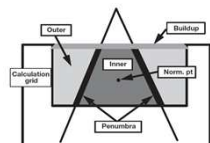


FIG. 11. Regions of different accuracy capabilities for photon beam dose calculations. Reproduced, with permission, from Ref. [10].

Ref: IAEA technique report 430

TABLE 18. EXAMPLE ILLUSTRATING DEVIATIONS (5) FOR DIFFERENT REGIONS
(Adapted, with permission, from Ref. [67].)

	Location	Type of region	1. Simple geometry (homogeneous)	2. Complex geometry (wedge, inhomogeneity, asymmetry)	More complex geometry (combinations of 1 and 2)
δ_1	Central beam axis	High dose, small dose gradient	2%	3%	4%
δ_2^*	Buildup region of central axis and penumbra regions of profiles	High dose, large dose gradient	2 mm or 10%	3 mm or 15%	3 mm or 15%
δ_3	Outside central beam axis	High dose, small dose gradient	3%	3%	4%
δ_4	Outside beam edges	Low dose, small dose gradient	3% ^a (30%)	4% ^a (40%)	5% ^a (50%)
RW_{rad}	Radiological width		2 mm or 1%	2 mm or 1%	2 mm or 1%
δ_{beam}	Beam fringe		2 mm or 3%	3 mm	3 mm

* These values are preferably expressed in mm. A shift of 1 mm corresponding to a dose variation of 5% is assumed to be a realistic value in the high dose, large dose gradient

^b This percentage is applicable to the following equation, $\delta_i = 100\% \times (D_{\text{calc}} - D_{\text{meas}}) / D_{\text{meas}}$, where $D_{\text{meas, cen}}$ is the dose on the central beam axis, since it is not always practicable to compare with the local dose. The values in brackets are those determined from Eq. (6).

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Example 1: basic photon test: open field

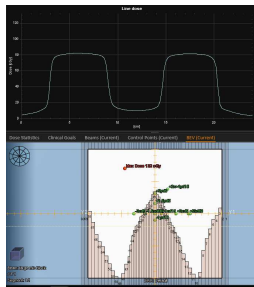
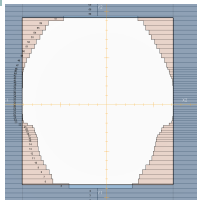
[illegible]

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Example 2 & 3: basic photon test: Small & large MLC-shaped field



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VALIDATION TESTS III- HETEROGENEITY CORRECTION VALIDATION

Confirmation of CT density table

Verifies dose beyond low-density (lung) material. The ratio of the dose values above and below the heterogeneous medium be measured and compared.



	Measured		planned	Measured vs. planned
10X	Mraw	Normalized dose (cGy)	(cGy)	% difference
point 1	22.8	115.94	115.5	0.38
point 2	15.26	77.60	77.1	0.65
Ratio		1.49	1.50	-0.26



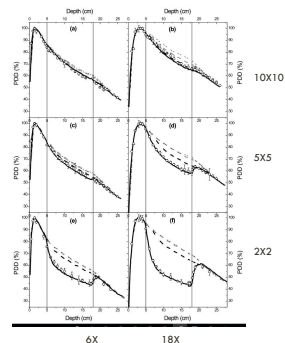
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The QMP must understand the limitations of the dose algorithms, particularly in the context of known dose discrepancies, which should be distinguished from incorrect commissioning of the TPS. Particular care should be taken when evaluating calculated dose 1) within low-density tissue, 2) near the interface of heterogeneous tissues, and 3) beyond low/high density tissue.



Ref: Carroasco et al. Med Phys. 2004;31(10):2899-911.



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VALIDATION TESTS IV- IMRT/VMAT

- Verification of small MLC field PDD
- Verification output for small MLC fields
- TG119 tests
- Clinical case tests
- RPC Phantom

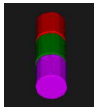


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TG119

Multitarget



Prostate



Head & Neck

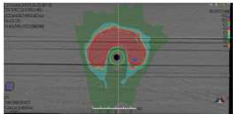


C Shape



Dose goals (anterior version)

Structure		
CShape PTV	95% of volume to receive at least 5000 cGy	10% of volume to receive no more than 5500cGy
Core	5% of volume to receive no more than 2500 cGy	



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TG119

Chamber measurements						
test	prescribed dose (Gy)	location	measured dose	planned dose	high dose region (mrts-plnt) perc.	low dose region (mrts-plnt) perc.
MultiTarget		isocenter				
		4 cm superior				
Prostate		4 cm inferior				
		isocenter				
Head/Neck		2.5 cm posterior				
		isocenter				
CShape (easy)		4.0 cm posterior				
		isocenter				
CShape (hard)		2.5 cm anterior				
		isocenter				
		2.5 cm anterior				
		isocenter				
		mean				
		standard deviation				
		confidence limit = (mean + 1.96 σ)				

Film measurements in phantom		
test	plane	% gamma pass
MultiTarget	isocenter	
	2.5 cm posterior	
Prostate	isocenter	
	4.0 cm posterior	
Head/Neck	isocenter	
	2.5 cm anterior	
CShape (easy)	isocenter	
	2.5 cm anterior	
CShape (hard)	isocenter	
	2.5 cm anterior	
mean		
standard deviation		
confidence limit = (100 - mean) ± 1.96 σ		

Field-by-Field % Gamma pass					
Field	MultiTarget	Prostate	Head/Neck	CShape (easy)	CShape (hard)
1					
2					
3					
4					
5					
6					
7					
8					
9					
mean					
overall mean					
standard deviation					
confidence limit = (100 - mean) ± 1.96 σ					



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Test patient: T6119

Case name: MultiTarget, Prostate, HeadNeck, Cshape

Test plan: IMRT GK

Ion Chamber/Array: MapCheck

Data acquired by: XL

Date: 3/15/2017

Note:

Field	MultiTarget (MF = 3.25)			Prostate (MF = 1.83)			HeadNeck (MF = 4.7)			Cshape (MF = 4.57)		
	3%/3mm	3%/2mm	2%/2mm	3%/3mm	3%/2mm	2%/2mm	3%/3mm	3%/2mm	2%/2mm	3%/3mm	3%/2mm	2%/2mm
1	100.0%	100.0%	99.3%	100.0%	100.0%	100.0%	99.5%	95.6%	88.0%	99.5%	97.6%	90.5%
2	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	99.5%	99.5%	97.8%	98.9%	94.7%	88.9%
3	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	97.7%	93.7%	100.0%	100.0%	100.0%
4	100.0%	99.4%	99.4%	100.0%	100.0%	99.1%	100.0%	100.0%	99.0%	100.0%	99.4%	99.4%
5	100.0%	100.0%	99.3%	100.0%	100.0%	100.0%	100.0%	100.0%	96.7%	99.5%	98.5%	89.2%
6	100.0%	98.7%	95.4%	100.0%	100.0%	98.5%	99.5%	98.9%	98.4%	100.0%	98.0%	92.5%
7	100.0%	99.3%	99.3%	100.0%	100.0%	100.0%	99.0%	97.0%	92.0%	100.0%	100.0%	98.7%
8							99.1%	97.3%	94.1%	100.0%	99.2%	98.4%
9							100.0%	98.4%	97.8%	99.5%	97.6%	93.2%
mean	100.0%	99.6%	99.0%	100.0%	100.0%	99.7%	99.6%	98.4%	95.2%	99.7%	98.4%	94.5%
Overall												
mean										99.8%	99.0%	96.8%
standard deviation										0.3%	1.4%	3.8%
confidence limit										0.8(99.2%)	1.7 (96.3%)	10.6(89.4%)

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CLINICAL VMAT CASES VALIDATION

Cases	3%/3mm	3%/2mm	2%/2mm
T spine (c-shape) 1	99.3%	97.5%	94.7%
T spine (c-shape) 2	99.3%	96.9%	92.3%
prostate+nodes 1	99.9%	99.3%	95.3%
prostate+nodes 2	100.0%	99.9%	98.1%
prostate+nodes 3	99.8%	99.6%	98.3%
prostate+nodes 4	100.0%	99.9%	99.8%
prostate+nodes 5	99.9%	99.4%	98.0%
H&N 1	99.8%	99.3%	97.7%
H&N 2	99.7%	96.9%	92.5%
H&N 3	99.7%	99.2%	96.3%
H&N 4	100.0%	99.7%	98.5%
H&N 5	99.6%	99.0%	93.2%
lung 1	100.0%	99.6%	98.4%
lung 2	99.4%	98.4%	97.5%
mean	99.7%	98.9%	96.5%
standard deviation	0.3%	1.1%	2.5%
confidence limit	0.9(99.1%)	3.3(96.7%)	8.4(91.6%)

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