A review of RTTQA audit

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CHART 8 hours on the machine





Chart cord

Table 5 Spinal cord dose, calculated vs. measured (mean doses (cGy))

Bronchus		Head and nec	k
Measured	Calculated	Measured	Calculated
37.6 ± 11.3 Difference: (Paired	34.3 ± 12.4 3.3 ± 1.3 p < 0.025	40.3 ± 20.7	37.3 ± 20.4 3.0 ± 1.4 p < 0.025)
mean ± SE)			

CHART QA: Phantoms(1st use of anatomical phantoms)

- Designed for treatments in Bronchus and Head and Neck (2D only)
- Outlines sent to centre in advance of visit
- Phantom set up by centre staff
- Dose delivered and measured promptly (using semi-flex-0.125cc- ion chamber) by visiting staff





CHART QA; some results of phantom measurements

- Dose delivered to the prescription point within 4% of 1.5Gy
- Variation of dose across volume: 5%
- Variation of dose to critical structures: very dependent on planning method......
- Dose to spinal cord lower in non-UK centres where only 2 fields (opposed pair) were used instead of 3 fields in UK
- Correction for lung: quite good! (most centres using stored data with bulk correction)



START QA Visit

- Measured dose to prescription point: average =0.985
- Cobalt60
- Incorrect normalisation point







Figure -1 Breast phantom showing measurement points

START Reference Point Chest wall phantom

Mean measured/expected dose 0.98

Tolerance 4%

Some centres implemented a 'lack of scatter correction'





Machine Issues found during audit visits - START

- Wedge
- Monitor ion chamber varying during the day
- Flatness at non zero gantry angles







Planning System A



Junction Phantom

 Overdoses of over 20% were found with some techniques in the junction between SCF and tangential fields



3D phantom (K Venables



Liz Miles)

Results: 14 planning systems Mean dose 0.987 (SD 0.013) All relative measurements within 5% of calculated; largest discrepancies at edge of field Small number of depts still not using lung correction





Figure 3-11 Accuracy of algorithms in the 3D breast phantom

PROSTATE PHANTOM for RT01 Moore AR, Warrington AP, Aird EG, Bidmead AM, Dearnaley DP

Constructed from water/WEP "Silver" prostate used for localisation Measuring points located in 3 planes

Small ion chamber for immediate dose measured at selected 3D points Independent Dose check with Alanine-from NPL







Slides into phantom - <u>before dosimetry</u> physical 1 sd ~ 1.6 mm 15 visits - no shift on phantom 2-3mm shift on phantom at 2 visits

All measured data differences from TPS



Some more RT01 results

- Rectal dose generally OK
-but plan data used critically to determine new Rectal Volume constraints (ref: Dose-volume constraints to reduce rectal side effects from prostate radiotherapy: evidence from MRC RT01 Trial ISRCTN 47772397.Gulliford SL, Foo K, Morgan RC, Aird EG, Bidmead AM, Critchley H, Evans PM, Gianolini S, Mayles WP, Moore AR, Sánchez-Nieto B, Partridge M, Sydes MR, Webb S, Dearnaley DP.

PARSPORT TRIAL TPS tests



* One centre failed due to film saturation in the high dose region.

Parsport Clark et al R+O 93(2009)102-109

Parsport

- 5-6 hours of machine measurement
- CIRS head and neck phantom
 - Conventional plan
 - IMRT plan

CONV IMRT	Average dose (Gy)	Dose range (Gy)	Centre n	Centre no.				Mean (%)	SD (%)	
			1 (%)	2 (%)	3 (%)	4 (%)	5 (%)	6 (%)		
PTV1	1.95 2.24	1.83–2.11 2.17–2.32	1.1 0.1	2.0 - 0.4	1.2 - 2.0	-0.9 1.8	0.0 - 2.4	0.9 - 0.5	0.7 - 0.6	1.0 1.5
PTV2	1.89 1.82	1.69–2.06 1.68–2.02	0.9 0.5	1.7 2.2	0.3 - 1.9	-0.3 - 2.7	-0.7 - 2.4	-2.6 - 0.7	-0.1 - 0.8	1.5 1.9
Cord	0.46 1.12	0.24-1.06 0.55-1.43	-1.7 - 0.5	0.0 2.4	-6.4	-35.8 4.2	<u>-32,7</u> - 4.9	2.9 - 2.6	-11,5 - 1,3	17.8 4.1

Combined point dose results for conventional (*italic*) and IMRT (bold) plans. Mean and standard deviation (SD) are of the data for the six centres. Negative results for percentage differences mean that the measured point was lower than the TPS calculated point.





Plan results (10 centres)

		II	MRT	Conventional		
	AIM	median	range	median	range	
PTV1 D ₉₅	> 61.8	61.8	61.2 - 63.2	47.5	36.0 - 60.5	
PTV2 D ₉₅	> 51.3	50.0	46.0 - 52.7	37.0	22.1 - 44.0	
SC _{max}	< 48.0	45.3	39.4 - 48.0	43.7	42.6 - 46.9	
CL parotid mean	< 24.0	26.3	22.0 - 29.4	63.9	56.1 - 65.9	
IL parotid mean	< 24.0	55.9	34.3 - 63.9	63.4	61.3 - 65.4	

All data in Gy

A national dosimetric audit of VMAT and Tomotherapy in the UK

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PTW Octavius II phantom with various detectors



3DTPS test plan



Tsang et al. Br J Radiol. 2013 Feb;86(1022)

Head and Neck plan measurements

Typical coronal and sagittal planes





Point dose differences in clinical plans





Gamma analysis results

	mean pass rate			
	Clinical	3DTPS		
2%/2mm	93.6%	91.5%		
3%/2mm	97.9%	96.3%		
3%/3mm	99.3%	98.3%		
4%/4mm	99.9%	99.7%		



Gamma analysis results

	mean pass	rate	percentage of planes >95% of γ<1		
	Clinical	3DTPS	Clinical	3DTPS	
2%/2mm	93.6%	91.5%	60.5%	56.1%	
3%/2mm	97.9%	96.3%	86.0%	75.0%	
3%/3mm	99.3%	98.3%	98.8%	88.6%	
4%/4mm	99.9%	99.7%	100%	100%	



Gamma analysis results at 2%/2mm

	mean pass rate	percentage of planes >95% of γ<1
Breast	99.8%	100%
Prostate and Nodes	94.9%	73.1%
Head and Neck	93.4%	55.4%
3DTPS	91.5%	56.1%



Rotational Audit Issues identified

- Lack of couch modelling
- Minimum leaf gap too small
- High modulation / high MUs
- Non-continuously variable dose rate
- Lack of information as to what some TPS/Linac combinations are capable of achieving
- Lasers and barometers



Conclusions

- A national dosimetry audit of rotational radiotherapy has been undertaken
- More than 93% of analysed planes achieved more than 95% pass rates for gamma parameters of 3%/3mm
- For many systems 3%/2mm were better criteria
- The majority of centres achieved accurate implementation of TPS modelling and delivery for VMAT and Tomotherapy
- Evaluation of the standards which others starting a VMAT program should be able to achieve



Conclusion

• The implementation of QA in radiotherapy has become vitally important in recent years. Often, as has been demonstrated here, a clinical trial has led the way to the general benefit of all patients receiving radiotherapy. By pursuing QA in the first year of the clinical trial, the standard of treatment was set and any later uncertainties when analysing the results were avoided. Wariness at each centre visited was replaced by active co-operation and satisfaction with the high standards that could be achieved and maintained. In addition, these visits gave an opportunity for mutual exchange of ideas.

Aird et al R+O 36(1995)235-245

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