

Conflicts of interest

Research support from Elekta Instruments, AB

Session Educational objectives

Identify weak points in the SRS/SBRT treatment process

Demonstrate how methodology described in TG-100 can be used to assess and prevent treatment events

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Part I: What could possibly go wrong?

What happens if you really mess up

The New York Times

THE RADIATION BOOM By WALT BOGDANICH

A Pinpoint Beam Strays Invisibly, Harming Instead of Healing Published: December 28, 2010

Radiation Offers New Cures, and Ways to Do Harm Published: January 23, 2010

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2009: 3 patients (+1 at another center) were overdosed during intracranial SRS treatments on a retrofitted linac. One patient deteriorated to near vegetative state.



http://medicalphysicsweb.org/cws/article/opinion/45334

2006-2007: 145 patients over-treated. Scatter factor measured with a farmer chamber at 20% of value measured with a pinpoint chamber.

2004-2009: 76 patients irradiated incorrectly from by a similar small field calibration error.

Events, except those which result from patient intervention, in which Dose differs from Rx (or dose that would result from Rx dose) by:

- > 5 rem (0.05 Sv) EDE, or
- > 50 rem (0.5 Sv) to organ or tissue, or
- > 50 rem shallow dose equivalent to skin

<u>Total dose</u> differs from Rx by >= 20%

Single Fx dose differs from Rx by >=50%

OR Dose given to wrong individual

OR

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AND

OR

Dose to tissue other than Tx site >50 rem (0.5 Sv) and >=50% of the dose expected from the administration defined in the Written Directive

NRC 10 CFR § 35.3045 Report and notification of a medical eventager SEAAPM 2017

Misadministrations and other terms

Reportable events	Misadministration			
atient or operator suffers a mechanical	X-ray teletherapy:			
jury	Wrong patient / wrong site			
/eekly teletherapy x-ray or electron dose iffers from planned by ≥15%	Calculated total dose differs from planned by >20% (>10% 3 or fewer fx)			
-ray brachytherapy dose differs from	Weekly dose differs > 30%			
lanned by ≥ 10%	X-ray brachytherapy:			
iagnostic x-ray exposure where uspected or actual long-term damage to	Wrong patient / wrong site			
rgans or systems occurs.	Total dose differs from planned by > 20%			
efinitions from				

More likely...this sort of thing can happen:

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Impact of target point deviations on control and complication probabilities in stereotactic radiosurgery of AVMs and metastases.

Treuer H, Kocher M, Hoevels M, et al., Radiother Oncol., 2006 Oct;81(1):25-32.



Kim et al., Inter- and intrafractional dose uncertainty in hypofractionated Gamma Knife radiosurgery, JACMP, 17(2), 2016.

If you practice SRS/SBRT long enough, you will make a mistake.

This isn't a talk about the gory details. (Sorry).

Here's where you can find the gory details....



Event Notification Reports

Event Notification Report (Last Month) (rew data text file) | Data Dictionary 🗏 (Microsoft Excel)
2010s: | 2016 | 2015 | 2014 | 2013 | 2012 | 2011 | 2010 |
2000s: | 2009 | 2008 | 2006 | 2007 | 2006 | 2005 | 2004 | 2003 | 2002 | 2001 | 2000 |

1990s: | 1999 |

http://www.nrc.gov/reading-rm/doc-collections/event-status/event/



http://wwwpub.iaea.org/MTCD/publications/PDF/Pub1084_web.pdf

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Part II: Why do things go wrong?

Radiosurgery is hard.

Uncertainty makes it harder.

People make it harder still.

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SRS/SBRT often has difficult constraints



Brain SRS: Pituitary adenoma (optic pathways within few mm)

High doses per fraction, small # fractions Fields that must conform to anatomy Inhomogeneous dose within tumor



Spine SRS: (spinal cord within few mm)

Sharp dose gradients outside target: 10%-25%/mm (GK) >10%/mm (linac)

Extremely high requirements for accuracy and precision!

Radiosurgery is hard.

Uncertainty makes it harder.

People make it harder still.

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A. Mack, H. Czempiel, H-J Kreiner, et. al., Med Phys 29(4), 2002

Procedural uncertainty = isodose uncertainty

What difference does a 10% change in dose make?



Frames do not have perfect immobilization ation!



SRS frames provide for low setup uncertainty and robust immobilization.

Practically limits treatment to single fraction.

Looks more invasive than it really is.

	Setup	Error						Intr	afraction	Error				
	Trans	lation	(mm)		Rotat	ion (°)		Tra	nslation	(mm)		Rotat	ion (°)	
	LR	AP	сс	Vector	LR	AP	CC	LR	AP	СС	Vector	LR	AP	CC
Mean	-0.19	0.08	-0.35	0.40	-0.14	-0.03	0.10	-0.03	-0.03	-0.03	0.05	-0.05	-0.03	-0.01
SD	0.32	0.29	0.50	0.66	0.25	0.19	0.20	0.05	0.18	0.12	0.22	0.30	0.20	0.09

Li, et al., IJROBP 2016.

...and neither do thermoplastic masks.



CyberKnife intrafraction motion vs time. n=50 patients. 2-mm thermoplastic mask immobilization. Thermoplastic masks are (possibly) more convenient than an SRS frame.

Tradeoff is masks have higher setup and intrafractional uncertainty.

Intrafraction shifts have been reported between 0.1 mm and 2.0 mm.

Mean 3D error (mm)

Require some kind of intrafraction motion management to achieve comparable performance to SRS frame.

C-W Wang, et al., Plos One, April 20, 2015

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...and neither do body frames!





Treatment site: technique (# patients/ # fractions)	Pretreatment	Intrafraction				
Lung: compressed (55/223))	7.29 ± 4.05	1.72 ± 1.98				
Lung: uncompressed (86/339)	7.40 ± 3.97	1.28 ± 1.53				
Liver (29/112)	6.64 ± 3.46	1.21 ± 1.74				
Prostate (48/240)	7.62 ± 3.97	1.95 ± 1.76				
Spine (45/91)	8.00 ± 4.43	1.29 ± 1.45				

R. Foster, et al., Localization Accuracy and Immobilization Effectiveness of a Stereotactic Body Frame for a Variety of Treatment Sites, IJROBP 87(5) 2013017

Machine calibrations are not always perfect







Percent frequency histogram of the RPC/INST ratios for the TG-51 non-compliant beams measured to date compared to the normal TG-51 compliant photon beams (n=600). With the exception of the Gamma Knife, the distribution of results for the TG-51 non-compliant beams. Gamma Knife shows a 4.5%, low result of which 1% is due to the muscle to water conversion and the remaining 3% due to differences between TG-21 in polystyrene and TG-1 in water dose calculations.

Quality Audits of the Calibration for TG-51 Non-Compliant Beams by the Radiological Physics Center

In some cases, accepted standards for output calibration do not exist (yet).

D. Followill, et al., Medical Physics 35:2774, 2008.

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Small fields are difficult to measure



Charged particle equilibrium assumption no longer valid

Detector itself becomes a prominent source of measurement uncertainty

Measured profile with different sized detectors

Ratio of absorbed doses vs field size for various detectors + monte-carlo

Volume averaging over detector makes field edge measurements inaccurate

Low, et al., Medical Physics 30(7), 2003. R. Alfonso, et al., Medical Physics 35(11), 2008.





Determination of the 4 mm Gamma Knife helmet relative output factor using a variety of detectors (summary of literature)

Note that ion-chamber measurement is well below other detectors.

JS Tsai et al., Med Phys 30(5), 2003.

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Not all planning systems work well for SRS/SBRT



J. Siebers, AAPM Summer School 2011 – Uncertainties in External Beam Radiation Therapy

TPS uncertainties

Beam data collection Beam data modeling MLC modeling Dose calc algorithms CT to election density Dose grid interpolation Accelerator output variations vs plan Range uncertainty (protons)

It can be difficult to decide on a target

Contours from 11 observers, 4 institutions



Delineation variability correlated with irregularity (r=0.77, p=0.005)

Peulen et al., Radiother Oncol, 114, 2015.



GTV_{3D} = free breathing CT

 $ITV_{MIP} = MIP$

ITV_{10phase} = combination of GTVs from each phase of 4DCT

 $ITV_{10phase}$ does not completely cover GTV_{3D} and ITV_{MIP}

Se et al., IJROBP 85, 2013.



Fig. 1. Contouring variations in a patient with a brainstem arteriovenous malformation. (a) Target volume presented as different colors. All contours were automatically projected on computed tomography images used for further analysis (b).

31 AVM patients 6 observers contouring on DSA Mean AR = 0.19 ± 0.14 AR never exceeded 0.6 in any patient!

Agreement ratio (AR) = $\frac{\text{common overlapping volume}}{\text{encompassing volume}}$

Stereotactic radiosurgery for brain AVMs: role of interobserver variation in target definition on digital subtraction angiography Buis, et al., IJROBP 62(1), 2005

Even subtle timing differences can matter

Timing of contrast injection can have significant effects on GTV definition

Immediate scan		tte scan Delayed scan					
Lesion Mean volume (SD) (mm ³)		Mean volume (SD) (mm ³) Mean volume (SD) (mm ³)			% change in volume	3D shift in isocentre (mm)	
A1	279	(79)	474	(59)	70	1.4	
A2	not analysed						
B1	290	(87)	325	(63)	12	1.0	
B2	879	(114)	1134	(103)	28	0.7	
С	477	(15)	492	(21)	3	0.9	
D1	1479	(32)	1798	(22)	21	1.3	
D2	1780	(33)	1767	(35)	-1	0.4	
E	1708	(33)	2093	(101)	22	0.6	
F1	1807	(21)	2731	(39)	51	0.5	
F2	2326	(23)	3179	(45)	36	1.5	
G1	1961	(161)	2871	(559)	46	0.2	
G2	3764	(234)	5952	(188)	58	1.4	
H1	5333	(138)	6434	(166)	20	2.4	
H2	not analysed						
I	11358	(344)	13047	(115)	14	4.6	
J	19787	(894)	16688	(5009)	-16	not planned	

Delayed scan was after a mean of 65 minutes Planners would select larger collimator sizes in 92% of delayed CT scans

Delineation of brain metastases on CT images for planning, radiosurgery: concerns regarding accuracy, K. Sidhu, P Cooper, R. Ramani, et. al. BR J Radiol (77), 2004. Schlesinger SEAAPM 2017



Scan 1: time of injection Scan 2: ~10 min delay Scan 3: ~15 min delay

Scans compared	% studies w ≥ 1 new lesion	95% CI	Range of # new lesions
Scan 1:2	35.3%	22.4%-49.9%	1-10
Scan 2:3	21.6%	11.3%-35.3%	1-9
Scan 1:3	43.1%	29.3%-57.8%	1-14

M. Kushnirsky et al., JNS 124, 2016







In in-vitro cultures, LQ model parameters overestimate BED as compared to empirical survival curves at SBRT doses

C. Park, L. Papiez, S. Zhang, M. Story, R.D. Timmerman, Int. J. Radiation Oncology Biol. Phys., 70(3), 2008

...so there is uncertainty in the LQ model

Observed in-vitro cell survival not as good as LQ model predicts

But clinically SRS performs better than LQ model predicts

Microvascular damage has been shown to occur at doses > 10Gy.

SRS biological effect may involve DNA damage + vascular damage



J. Kirkpatric, J. Meyer, L.B. Marks, Semin Rad Onc, 18(4), 2008.

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You can't localize targets perfectly



Modality	Radial Deviation (mean ± STD) (mm)
СТ	0.4±0.2
*MR (T1-weighted)	1.4±0.3
*MR (T2-weighted)	1.4±0.5
[†] PET	1.1±0.5
[‡] SPECT	1.6±0.5

*Siemens Magnetom Symphony, †Siemens CTI ECAT EXACT HR, ‡Siemens MULTISPECT 3

Localization of known stereotactic targets with various modalities

C.P. Karger, P. Hipp, M. Henze, G. Echner, et al., Phys Med Biol 48, 2003.

Internal anatomy moves!





Be careful of dose interplay effects!

Images courtesy of S. Benedict, UC-Davis

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...and images are not necessarily reality!





Radiosurgery is hard.

Uncertainty makes it harder.

People make it harder still.

Humans are (maybe) not so good at 3D registration



Humans are (maybe) not so good at 3D registration



Can you spot the differences between these two registrations?

Humans are (maybe) not so good at 3D registration



 Δ translation x: -1.00 mm Δ translation y: -0.64 mm Δ translation z: 1.52 mm

 Δ rotation x: 0.05° Δ rotation y: -2.00° Δ rotation z: 0.00°

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Humans are (maybe) not so good at 3D registration



 Δ translation x: -1.00 mm Δ translation y: -0.64 mm Δ translation z: 1.52 mm Δ rotation x: 0.05° Δ rotation y: -2.00° Δ rotation z: 0.00°

Even "simple" procedures can go wrong



Chronic rib fracture/fragmentation post SBRT

Images courtesy of University of Virginia

Frame pin pushed through skull defect Schlesinger SEAAPM 2017

SRS/SBRT can have unusual machine settings



Couch 0.0° / Gantry 65.3° No collision



Couch 323.9° / Gantry 65.3° Collision!

Beams are often non-coplanar, requiring unusual couch angles. Not difficult to have a collision with patient!

Image from D. Schlesinger, et al., Treatment Planning for Spine Radiosurgery, in Spine Radiosurgery 2nd ed., 2015.

SRS/SBRT treatment plans require extreme care



Clinical plan with narrow beam arrangement and no accounting for couch or immobilization devices



Replan including 1 cm bolus to simulate couch and immobilization

Actual grade 4 skin necrosis



Hoppe et a., IJROBP 72. 2008.

Variations in technique can be large



mean dose to PTV: $99.7 \pm 3.5\%$ mean dose to V98%: $93.6 \pm 4.4\%$, respectively

Esposito et al., Physica Medica 32 (2016)



14 centers, 5 patient image sets w single liver metastasis Common set of dose constraints Local equipment, planning technique

One theme so far is that humans are fallible!

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Part III: How not to be a victim



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Prescriptive QA for SRS/SBRT

AAPM Task Group	Title	Year Published
TG-42	Stereotactic Radiosurgery	1995
TG-53	QA for clinical radiotherapy treatment planning	1998
TG-51	Clinical reference dosimetry of high-energy photon and elect on beems	1999
TG-142	QA of medial accelerators	2009
TG-101	Sterepta the budy radiation therapy	2010
TG-148	Q for hencal tomotherapy	2010
TG-135	QA for robotic radiosurgery	2011
TG-179	QA for image-guided radiation therapy using CT- based technologies	2012
TG-147	QA for nonradiographic radiotherapy localization and positioning systems	2012 Schlasinger SEAADM 2017

	Exa	am	nple: T	G-	14	2 (A	ACCE	eler	ator	QA)	
TABLE III.	. Annual.						TABLE VL Imaging.				
			Machine-type tolerance							Application-type tolera	nce
Procedure		Non-IMRT	IMRT	SRS/SI	BRT		Procedure		no	n-SRS/SBRT	SRS/SBRT
Dosimetry	у								Daile		
X-ray flat X-ray syn Electron f Electron s SRS arc r (range:	ness change from baseline nmetry change from baseline flatness change from baseline symmetry change from baseline outation mode 0.5–10 MU/deg)	NA	1% ±1% 1% ±1% NA	Monitor units set 0 MU or 2% (whi	vs delivered: hever is greater)	TABLE II. Monthly.	Planar kV and MV (EPII Collision interlocks Positioning/repositioning	D) imaging	Dany	Functional ≤2 mm	Functional ≤1 mm ≤1 mm
				Gantry arc set 1.0° or 2% (which	ever is greater)				Machine-type tolerar	ice	
X-ray/elec Spot chec output f (two or	ctron output calibration (TG-51) & of field size dependent lactors for x ray more FSs) torse of electron applicators		±1% (absolute) 2% for field size <4×4 cm ² , 1% ≥4×4 cm ² +2% form bandling			Procedure Desimetry		Non-IMRT	IMRT	SRS/SBRT	Functional ≤1 mm ≤1 mm
(spot ch X-ray bea Electron b	eck of one applicator/energy) m quality (PDD ₁₀ or TMR ³⁰) beam quality (R ₅₀)		±1% from baseline ±1 mm			Electron output constancy Backup monitor chambe Typical dose rate ⁸ output	y r constancy t constancy	NA	2% 2% (@ IMRT dose rate)	2% (@ stereo dose rate, MU)	<1 mm
Finysteal of factor of X-ray mo (output) Electron r	enge transmission enstancy constancy) monitor unit linearity	±2% ≥5 MU	±2% ±5% (2-4 MU), ±2% ≥5 MU ±2% ≥5 MU	±5% (2-4 ±2% ≥5	MU), MU	Photon beam profile con Electron beam profile co Electron beam energy co	istancy instancy instancy		1% 1% 2%/2 mm		≤2 mm Baseline Baseline
X-ray ou X-ray ou Electron	TABLE I. Daily.					Mechanical Light/radiation field coin Light/radiation field coin	acidence ^b acidence ^b (asymmetric)		2 mm or 1% on a side 1 mm or 1% on a side		Baseline ≤1 mm
gantry a Electron			Machine	-type tolerance		front pointer	or lasers compared with		Imm		
constan Arc mode	Procedure		Non-IMRT	IMRT	SRS/SBRT	Gantry/collimator angle (@ cardinal angles) (d	indicators igital only)		1.0°		≤1 mm Baseline Baseline
TBI/TSE PDD or 1 TBI/TSE TBI/TSE	Dosimetry X-ray output constancy (all energies)				Jaw position indicators (Jaw position indicators (Jaw position indicators (Crows bair contering (we	(symmetric) ^e (asymmetric) ^d		2 mm 2 mm 1 mm		Baseline
Mechani Collimate Gantry re Couch ro	Electron output constance except for machines wi e-monitoring requiring	y (weekly, ith unique daily)		3%		Treatment couch positio Wedge placement accur Compensator placement Latching of wedges, blo Localizing lasers	nindicators ^e ncy accuracy ^f cking tray ^g	2 mm/1* ±2 mm	2 mm/1° 2 mm 1 mm Functional ±1 mm	1 mm/0.5° < ± 1 mm	≤1 mm Baseline Baseline Baseline Baseline
Coincider mechan Table top Table ang Table tra	Laser localization Distance indicator (ODI) Collimator size indicator	@ iso	2 mm 2 mm 2 mm	1.5 mm 2 mm 2 mm	1 mm 2 mm 1 mm	Safety Laser guard-interlock ter Respiratory gating	si		Functional		
movem Stereotac	Safety Door interlock (beam off	D	P	unctional				E. Klein	, et al., Mee	d. Ribyrs 36(2)	PH2009.





WL pointer with embedded BB at end



WL pointer mounted to treatment couch

Winston-Lutz test

Determines alignment of mechanical, radiation, and imaging isocenters of a linac

> Mount WL-pointer with embedded BB to the end of the treatment couch (usually special mounting hardware)

Align BB end of pointer to the isocenter as suggested by the room lasers

MV and KV detectors used to ensure isocenter alignment

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Regular QA of imaging systems is critical for SRS/SBRT! This includes after hardware/software upgrades.

SRS/SBRT end to end tests





Tests the entire treatment procedure

Remember that phantoms provide a best case measure of uncertainty









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Patient-specific QA

Performed using phantoms that have arrays of ion chambers or diodes Assures that machine can technically deliver a given treatment plan











- 1. Recalculate plan onto image of phantom
- 2. Deliver treatment to phantom
- 3. Analyze results

Be careful! Not patient geometry!

Evaluating patient-specific QA

Popular metric: Gamma Index



Be careful: How to choose criteria?

Dose difference: Percentage difference on a pixel by pixel basis.

Distance to agreement (DTA): Distance to closest point with same dose

Gamma index looks for pixels where dose difference and DTA are simultaneously greater than a pre-selected threshold (example: 1 mm/1% or 3mm/3%).

Results usually expressed as a passing rate (example: >95% passing)

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SRS detectors are critical!



4x4 cm2 field, cross-beam profile

CC13 ion chamber vs diode vs deconvolved ion chamber signal

Ion chamber: 0.13 cm2 active volume

Diode: 0.8x0.8 mm2 cross sectional area

Use dedicated stereotactic ion chambers or diodes for making high-resolution measurements!

The extraction of true profiles for TPS commissioning and its impact on IMRT patient-specific QA, Yan, et al., Med Phys 35(8), 2008 _{Schlesinger SEAAPM 20}



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Apply consensus experience

TARLE III. Summary of suggested dose constraints for various critical organs. Note that for serial tissues, the volume-dose constraints are given in terms of the critical maximum tissue volume that should receive a dose equal or greater than the indicated threshold dose for the given number of fractions used. For parallel tissue, the volume-dose constraints are based on a critical minimum volume of tissue that should receive a dose equal to rels shan the indicated threshold dose for the given number of fractions used.

		One f	raction	Three f	ractions	Five fr	actions	
Serial tissue	Max critical volume above threshold	Threshold dose (Gy)	Max point dose (Gy) ^a	Threshold dose (Gy)	Max point dose (Gy) ^a	Threshold dose (Gy)	Max point dose (Gy) ^a	End point (≥Grade3)
Optic pathway	<0.2 cc	8	10	15.3 (5.1 Gy/fx)	17.4 (5.8 Gy/fx)	23 (4.6 Gy/fx)	25 (5 Gy/fx)	Neuritis Hearing
Cochlea			9		17.1 (5.7 Gy/fx)		25 (5 Gy/fx)	loss
Brainstem								Cranial
(not medulla)	<0.5 cc	10	15	18 (6 Gy/fx)	23.1 (7.7 Gy/fx)	23 (4.6 Gy/fx)	31 (6.2 Gy/fx)	neuropathy
Spinal cord	<0.35 cc	10	14	18 (6 Gy/fx)	21.9 (7.3 Gy/fx)	23 (4.6 Gy/fx)	30 (6 Gy/fx)	Myelitis
and medulla	<1.2 cc	7		12.3 (4.1 Gy/fx)		14.5 (2.9 Gy/fx)		
Spinal cord subvolume								
5-6 mm above	<10%							
and below level	of							
reated per Ryu)	subvolume	10	14	18 (6 Gy/fx)	21.9 (7.3 Gy/fx)	23 (4.6 Gy/fx)	30 (6 Gy/fx)	Myelitis
Cauda equina	<5 cc	14	16	21.9 (7.3 Gy/fx)	24 (8 Gy/fx)	30 (6 Gy/fx)	32 (6.4 Gy/fx)	Neuritis
Sacral plexus	<5 cc	14.4	16	22.5 (7.5 Gy/fx)	24 (8 Gy/fx)	30 (6 Gy/fx)	32 (6.4 Gy/fx)	Neuropathy
Esophagus ^b	<5 cc	11.9	15.4	17.7 (5.9 Gy/fx)	25.2 (8.4 Gy/fx)	19.5 (3.9 Gy/fx)	35 (7 Gy/fx)	Stenosis/fistula
Brachial plexus	<3 cc	14	17.5	20.4 (6.8 Gy/fx)	24 (8 Gy/fx)	27 (5.4 Gy/fx)	30.5 (6.1 Gy/fx)	Neuropathy

Organ at risk tolerance data is sparse

Good references: AAPM TG-101, RTOG-0915, RTOG-0236, RTOG-0618, RTOG-1112, RTOG-0813, QUANTEC

If you experiment, formalize as an approved clinical trial! Benedict, et al., AAPM Task Group 101, Medical Physics, 37(8), 2010.



Potters, et al., ASTRO and ACR Practice Guidelines for the Performance of Stereotactic Body Radiation Therapy, IJROBP 76(2), 2010.

Solberg, et al., Quality and Safety Considerations in Stereotactic Radiosurgery and Stereotactic Body Radiation Therapy, PRO, Aug 2011.

Benedict, et al., Stereotactic Body Radiation Therapy: The report of AAPM Task Group 101, Medical Physics, 37(8), 2010.

Perform QA specific for SRS/SBRT

Follow accepted best practices

Value training, credentialing, and peerreview

Apply systems engineering

Keep innovating

Credentialing and training

Quality and Safety Considerations in Stereotactic Radiosurgery and Stereotactic Body Radiation Therapy

Timothy D. Solberg, Ph.D.¹, James M. Balter, Ph.D.², Stanley H. Benedict, Ph.D.³, Benedick A. Fraass, Ph.D.2, Brian Kavanagh, M.D.⁴, Curtis Miyamoto, M.D.⁵, Todd Pawlicki, Ph.D.⁶, Louis Potters, M.D.⁷, Yoshiya Yamada, M.D.⁸

"If the radiation oncologist's formal training did not include SRS/SBRT, then specific training in SRS/SBRT, including a minimum of 5 CME credit hours and direct observation of treatment of at least 3 different patients, should be obtained prior to performing any SRS or SBRT procedures"

Same idea for medical physicist and neurosurgeon!



Include charts, QA, clinical procedures – anything and everything!



Treatment machines are just like manufacturing machines



Kuka 240-2 Robot (Kuka Roboter, GMBH, Augsberg)



CyberKnife G4 (Accuray, Sunnyvale)

So it is logical to look to systems engineering for guidance!



Can we even do sub-mm QA for SRS/SBRT?

Device	Detector spacing	
Mancheck 2	70 mm	
ArcCheck	10.0 mm (effective 7.0 mm w helical geometry)	
Octavius 1000 SRS	2.5 – 5.0 mm	
MatriXX	7.6 mm	
SRS MapCheck (2017 release)	2.5 mm	
Radiochromic film	Limited by scanner resolution / uncertainty! Ima	age courtesy of Sonja Dieterich, Ph.D.
Doco gradiant of 16 Cv/	mm > challonging over fo	r filml

Dose gradient of 16 Gy/ mm -> challenging even for film!



Hierarchy of mistake-proofing principles (poka-yoke), various sources (usually attributed to Shigeo Shingo and the Toyota Production Method)

Systems engineering tools

Incident Learning: Lessons from your (almost) mistakes

Statistical Process Control: Detect changes in your process

Process Mapping: Think through and diagram clinical process

Failure Mode and Effects Analysis (FMEA): start with process, identify and prioritize failure modes

Fault Tree Analysis: Start with failure mode, identify causes and find opportunities to improve

Report and learn from almost mistakes

METRIC	AGGREGATE CURRENT QUARTER	AGGREGATE HISTORICAL SUM
Reported Events	274	2345
Therapeutic Radiation Incidents	58	645
Near Miss	79	773
Unsafe Conditions	89	695
Operational/Process Improvement	27	61
Most Commonly Identified Workflow Step Where Event Occurred	Treatment Planning: 30% (83/274)	Treatment Planning: 28% (662/2345)
Most Commonly identified Workflow Step Where Event was <i>Discovered</i>	Treatment Delivery Including Imaging (e.g. at the machine): 28% (77/274)	Pre-treatment QA Review (e.g. Physics Plan Check): 25% (580/2345)
Most Commonly Identified Treatment Technique	3∙D: 27% (101/274)	3-D ; 21% (514/2345)
Treatment Technique	27% (101/274)	21% (514/2345)

https://www.astro.org/RO-ILS-Education.aspx

Incident learning systems

Report mistakes and almost mistakes

Focus on process improvement, not assigning blame

Foster a sense that group learning is of critical importance

RADIATION ONCOLOG INCIDENT LEARNING SYSTEM Sponsored by ASTRO and AAPM





Machine logs are analyzed for planned vs actual machine parameters (such as MLC positions and beam on/off status).

Fluence maps are reconstructed based on actual machine parameters during delivery.

Comparisons made against planned fluence maps.

Method can overcome limitations of phantombased patient-specific QA. ..and logfilebased QA



Handsfield, et. al., Med Phys 41(10), 2014

Perform QA specific for SRS/SBRT

Follow accepted best practices

Value training, credentialing, and peerreview

Apply systems engineering

Keep innovating



The future: Autosegmentation

De novo, segmented edit, peer and self-edit

Segmented edits remained closest to ground truth

Segmentation editing improves efficiency while reducing inter-expert variation and maintaining accuracy for normal brain tissues in the presence of space-occupying lesions M.A. Deeley, et al., Phys Med Bio 58 (2013)



A1 while the other colors represent manual expert segmentation

red contours are those of the

The future: Updated RadBio modeling

Author/Year	Model Name	Strategy
Guerrero and Allen (2004)	Modified LQ model (MLQ)	Linear-Quadratic-Linear
Park, et al. (2008)	Universal Survival Curve	Hybrid LQ and multi-target model
Kavanagh and Newman (2008)	Kavanagh- Newman	Dose-dependent increase in exponential rate of cell kill
Astrahan (2008)	L-QL model	Linear-quadratic linear
Hanin and Zaider (2010)		Microdosimetry model
Wang, et. al. (2010)	Generalized Linear Quadratic Model (gLQ)	Adds a parallel β_2 term to account for less sub-lethal repair at high doses



The future: Treatment planning automation?



Software automatically segments brain metastases and creates radiosurgeryready plans using multiple conformal arcs. Treatment planning takes minutes. Removes the human component (and error?).

T. Gevaert, et al., Radiation Oncology (11) 2016

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In the next presentation:

Systems engineering tools

Incident Learning: Lessons from your (almost) mistakes

Statistical Process Control: Detect changes in your process

Process Mapping: Think through and diagram clinical process

Failure Mode and Effects Analysis (FMEA): start with process, identify and prioritize failure modes

Fault Tree Analysis: Start with failure mode, identify causes and find opportunities to improve

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Conclusions

SRS/SBRT are complicated procedures with many sources of uncertainty.

There are many ways to have a misadventure.

There are proven ways to reduce risk:

Training and credentialing Formal analysis of procedural risk Constant learning – including close calls Formalizing new techniques as clinical trials



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Department of Radiation Oncology