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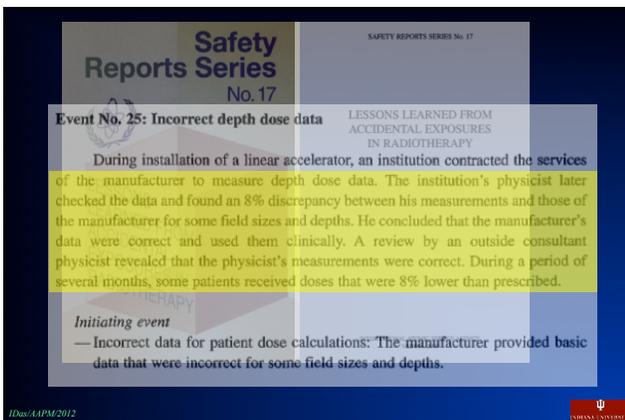
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## Planning for Commissioning Time

$$\text{Time} = \frac{(\text{PDD} + 5 \text{ profiles})}{\text{beam energy}} \times (\text{open} + 4 \text{ wedges}) \times 60 \text{ points/scan} \times [(1 \text{ s/pts} + (1 \text{ s/movement and delay})) \times (15 \text{ fields} \times 2 \text{ energies})]$$

~10<sup>5</sup> s  
~ 30 h

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## Rational For Commissioning Beam Data

The following concerns should be carefully evaluated before the use of any golden beam data within a clinic. First, it is not evident that manufacturing procedures for all linear accelerators have produced a level of reproducibility acceptable for clinical use. For example, variations in beam parameters have been noted between beams with the same nominal energies.<sup>3-5</sup> Second, on-site changes made during installation and acceptance of the user's accelerator (e.g., changes in beam energy and/or profiles from beam steering) will not be modeled in the golden data. Third, the beam characteristics of the soft wedges are made by moving jaws that depend on the speed parameters of the jaws and a deviation at site could affect the beam profile of the soft wedge. Fourth, although acceptable agreement with the golden data set may be found in individual checks, it may be that some clinical setups will have multiple errors, which combine to produce unacceptable results. Finally, the commissioned beam data also provide a thorough check of the accelerator, which may uncover problems that may not otherwise be discovered with a mere spot check.

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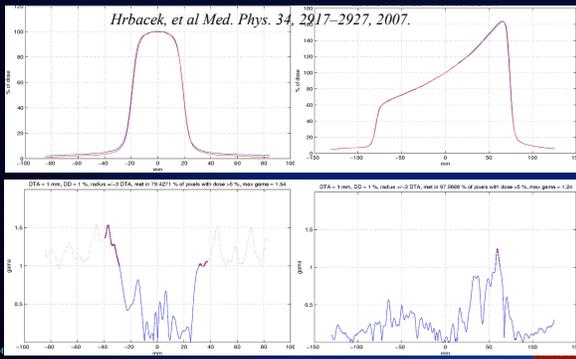
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## Rational For Not Using Golden Data



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### Rational For Not Using Golden Data

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$$|x_i - x^-| < \Delta, \forall x_i$$

$\Delta = ?$  (0.5, 1.0 Or 2%)

For all practical purposes, based on the presented results, we suggest 2 mm DTA and 2% DD as a convenient criteria for  $\gamma$  analysis to be met when evaluating the agreement of profiles scanned in common dosimetrical conditions. Better results are attainable by employing different strategies coping with the imperfections of measurements.

It is our opinion that matched beams which do not meet the earlier suggested criteria should not be treated as clinically interchangeable.

*Hrbacek, et al "Quantitative evaluation of a beam-matching procedure using one dimensional gamma analysis," Med. Phys. 34, 2917–2927, 2007.*

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Medical Physics, 39(2), 569-572, 2012

#### POINT/COUNTERPOINT

*Suggestions for topics suitable for these Point/Counterpoint debates should be addressed to Colin G. Orton, Professor Emeritus, Wayne State University, Detroit: oronc@comcast.net. Persons participating in Point/Counterpoint discussions are selected for their knowledge and communicative skill. Their positions for or against a proposition may or may not reflect their personal opinions or the positions of their employers.*

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**Vendor provided machine data should never be used as a substitute for fully commissioning a linear accelerator**

Indra J. Das, Ph.D.  
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 (Tel: 317-944-1303; E-mail: idas@iupui.edu)*

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Colin G. Orton, Ph.D., Moderator  
 (Received 22 September 2011; accepted for publication 25 September 2011; published 11 January 2012)  
 [DOI: 10.1118/1.3658740]

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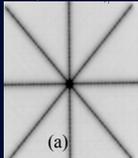
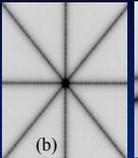
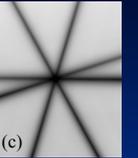
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#### Question

Star shot provides isocentricity of accelerator parameters. Which one is gantry star shot?

(a)      (b)      (c)

- 2% A. (a)
- 2% B. (b)
- 79%** C. (c)
- 14% D. (a) or (b)
- 4% E. None

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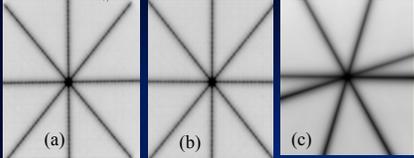
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Question  
Star shot provides isocentricity of accelerator parameters. Which one is gantry star shot?



A. (a)  
B. (b)  
C. (c)  
D. (a) or (b)  
E. None

Answer: C

Reference: Khan, Physics of Radiation Therapy, 2009

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Question  
Time required for commissioning a dual energy linear accelerator with photon and electron beam is

0% A. 1 day  
5% B. 3 days  
42% C. 1 week  
44% D. 4-6 weeks  
9% E. 2 months

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Question  
Time required for commissioning a dual energy linear accelerator with photon and electron beam is

A. 1 day  
B. 3 days  
C. 1 week  
D. 4-6 weeks  
E. 2 months

Answer: D

Reference: Das et al, TG-106, Med. Phys. 35(9), 4186-4214, 2008

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### Definition of Detectors

- ❖ **Standard chamber**  $10^{-1} \text{ cm}^3$ —The active volume for a standard Farmer-type ionization chamber is on average  $0.6 \text{ cm}^3$ .
- ❖ **Minichamber**  $10^{-2} \text{ cm}^3$ —The active volume for a mini-ionization chamber is on average  $0.05 \text{ cm}^3$ .
- ❖ **Microchamber**  $10^{-3} \text{ cm}^3$ —The active volume for a microionization chamber is on average  $0.007 \text{ cm}^3$  and ideally suited for small field dosimetry such as radiosurgery, gamma knife, CyberKnife, and IMRT

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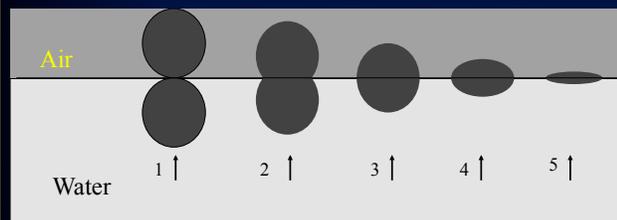
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### Setting Water Tank & Detector



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### Know Your Connectors



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### Understand Detector, Connector & Cable



Srivastava et al, SU-GG-T-270, 2010

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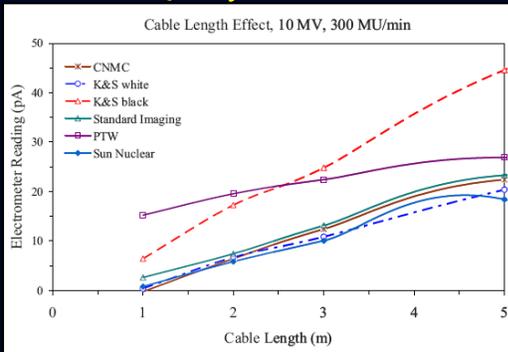
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### Quality of Cables



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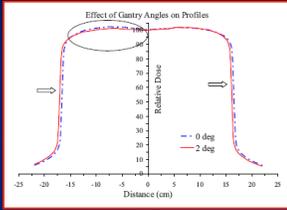
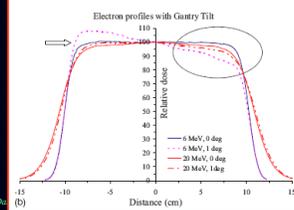
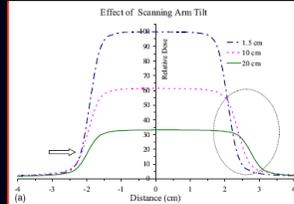
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### Setup and Possible Errors



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## Electrometer

- ❖ Null Setting
- ❖ Cable subtraction
- ❖ Proper bias
  - ⊠ >300 V for ion chamber
  - ⊠ 100 V for diamond
  - ⊠ 0 v for all diodes
- ❖ Proper gain
- ❖ Proper mode

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Question

### High voltage is applied to ion chamber for?

(1) To reduce ion recombination; (2) To reduce polarity effect; (3) 100 volts;  
(4) 300-400 volts

<b>15%</b>	<b>A.</b>	1 only
<b>15%</b>	<b>B.</b>	1 and 2 only
<b>0%</b>	<b>C.</b>	1, 2 and 3 only
<b>11%</b>	<b>D.</b>	2 and 3 only
<b>59%</b>	<b>E.</b>	1 and 4 only

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Question

### High voltage is applied to ion chamber for?

(1) To reduce ion recombination; (2) To reduce polarity effect; (3) 100 volts;  
(4) 300-400 volts

- A. 1 only
- B. 1 and 2 only
- C. 1, 2 and 3 only
- D. 2 and 3 only
- E. 1 and 4 only

**Answer: E**

Reference: Das et al, TG-106, Med. Phys. 35(9), 4186-4214, 2008

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**Question**

When setting ion chamber in water tank, the correct position of the chamber as viewed in water tank (as seen in figure) is:

2% 1. Position # 1  
 0% 2. Position # 2  
 89% 3. Position # 3  
 5% 4. Position # 4  
 4% 5. Position # 5

IBAS/AAP Reference: Das et al, TG-106, Med. Phys. 35(9), 4186-4214, 2008

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**Question**

When setting ion chamber in water tank, the correct position of the chamber as viewed in water tank (as seen in figure) is:

A. Position # 1  
 B. Position # 2  
 C. Position # 3  
 D. Position # 4  
 E. Position # 5

**Answer: C**

IBAS/AAP Reference: Das et al, TG-106, Med. Phys. 35(9), 4186-4214, 2008

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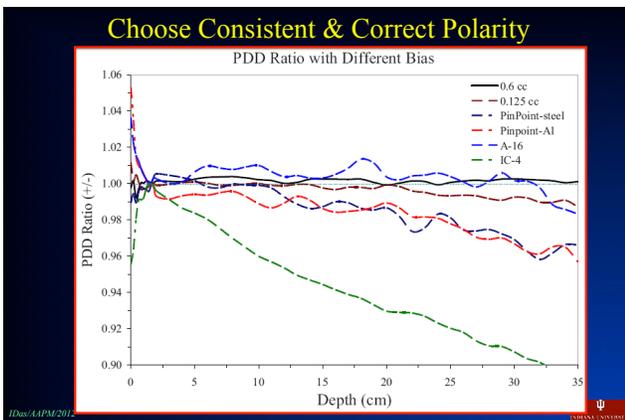
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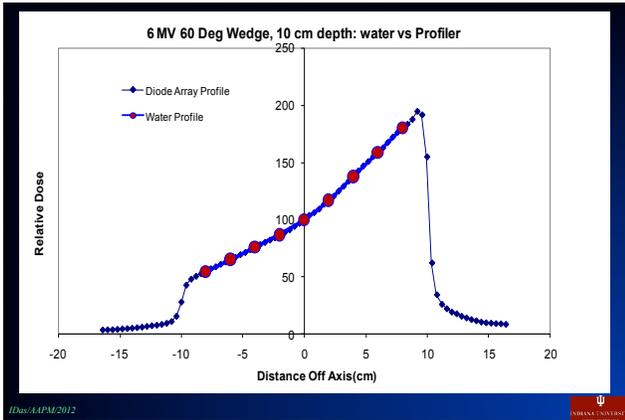
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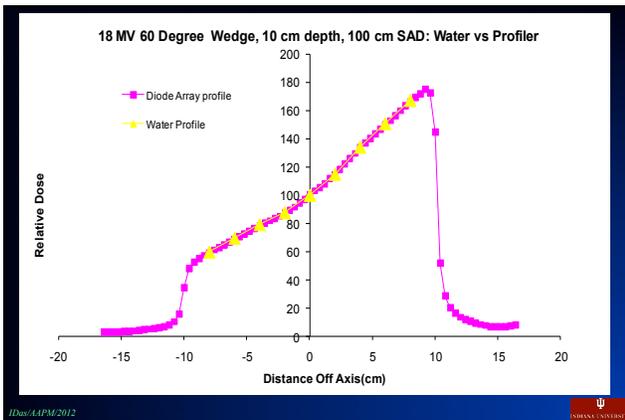
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**Question**

The possible setup error that causes the photon beam dose profile in figure is due to:

- 25% A. Gantry tilt
- 0% B. Collimator rotation
- 19% C. Tank arm tilt
- 26% D. Gantry and arm tilt
- 30% E. Gantry tilt, collimator rotation and tank arm tilt

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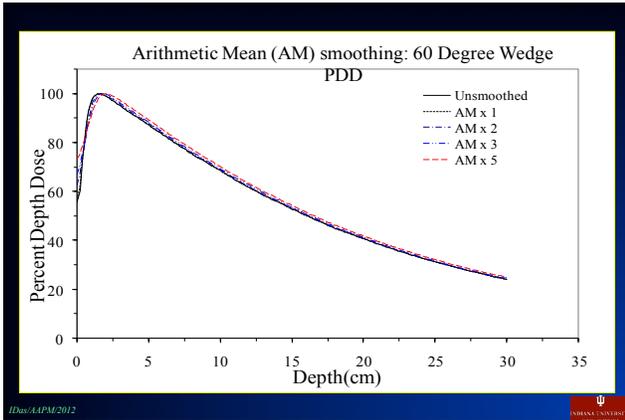
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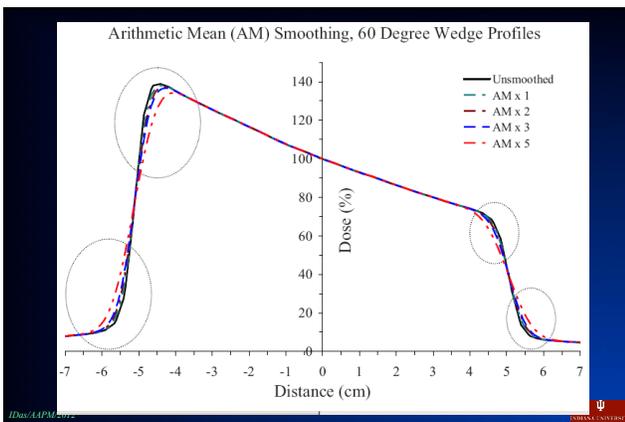
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## Future of Beam Data Commissioning

- ❖ Standardization of linear accelerators
- ❖ Monte Carlo based commissioning
- ❖ Newer Radiation Detectors & Cables
- ❖ Newer Scanning Systems
- ❖ Smart algorithms

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**An integrated 6 MV linear accelerator model from electron gun to dose in a water tank**

J. St. Aubin  
 Department of Physics, University of Alberta, 11322-89 Avenue, Edmonton, Alberta T6G 2G7, Canada  
 and Department of Oncology, Medical Physics Division, University of Alberta, 11560 University Avenue,  
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B. G. Fallone<sup>1)</sup>  
 Department of Physics, University of Alberta, 11322-89 Avenue, Edmonton, Alberta T6G 2G7, Canada;  
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 11560 University Avenue, Edmonton, Alberta T6G 1Z2, Canada

(Received 9 September 2009; revised 18 March 2010; accepted for publication 22 March 2010;  
 published 28 April 2010)

*Aubin et al., Med Phys, 37(5), 2279-2288, 2010*

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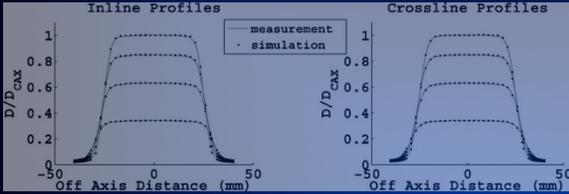
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**Simulated Profiles**



*Aubin et al., Med Phys, 37(5), 2279-2288, 2010*

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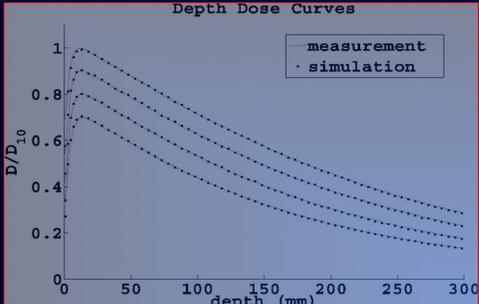
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**Depth Dose Simulation**



*Aubin et al., Med Phys, 37(5), 2279-2288, 2010*

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Question

Electron beam depth doses shown in figure represents the problem of

- 4% 1. Noise in the cable
- 2% 2. Electrometer gain
- 1% 3. Bias on the electrometer
- 91% 4. Speed of scanning
- 0% 5. Tuning of accelerator

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Question

Electron beam depth doses shown in figure represents the problem of

- A. Noise in the cable
- B. Electrometer gain
- C. Bias on the electrometer
- D. Speed of scanning
- E. Tuning of accelerator

Answer: D

Reference: Das et al. TG-106, Med. Phys. 35(9), 4186-4214, 2008

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Question

Photon beam dose profiles taken with various detectors as shown in figure is possibly due to:

- 4% A. Speed of scanning
- 0% B. Beam asymmetry
- 0% C. Pb piece in the beam
- 4% D. Hysteresis of scanning system
- 97% E. Orientation of scanning detector

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**Question**

Photon beam dose profiles taken with various detectors as shown in figure is possibly due to:

- A. Speed of scanning
- B. Beam asymmetry
- C. Pb piece in the beam
- D. Hysteresis of scanning system
- E. Orientation of scanning detector

**Answer: E**

Reference: Das et al, TG-106, Med. Phys. 35(9), 4186-4214, 2008

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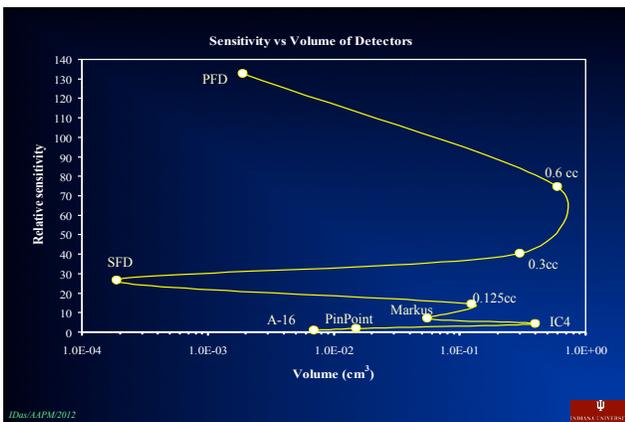
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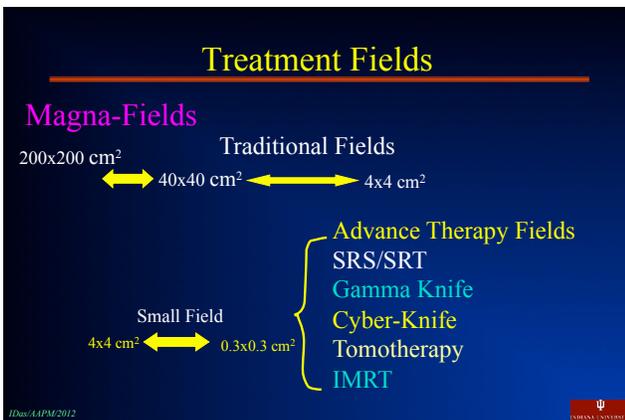
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## What is a Small Field?

- ❖ Lack of charged particle
  - ▣ Dependent on the range of secondary electrons
  - ▣ Photon energy
- ❖ Collimator setting that obstructs the source size
- ❖ Detector is comparable to the field size

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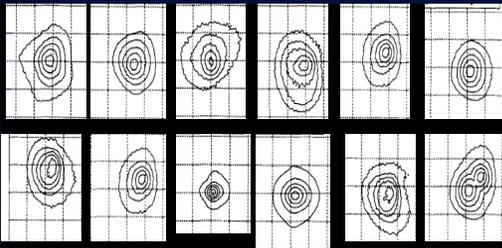
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## Views of Source Sizes



Jaffray et al, Med Phys 20, 1417-1427 (1993).

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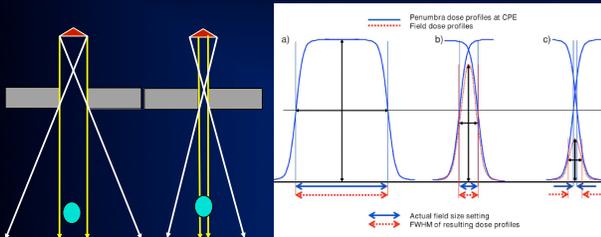
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## Definition of Small Fields



Das et al, Med. Phys. 35, 206-215, 2008

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## Dosimetry

- ❖ Absolute
  - ⌘ Dose
- ❖ Relative
  - ⌘ Depth Dose  $[D(r,d)/D(r,d_m)]$
  - ⌘ TMR
  - ⌘ Profiles
  - ⌘ Output,  $S_{cp}$  (total scatter factor),  $[D(r)/D(ref)]$

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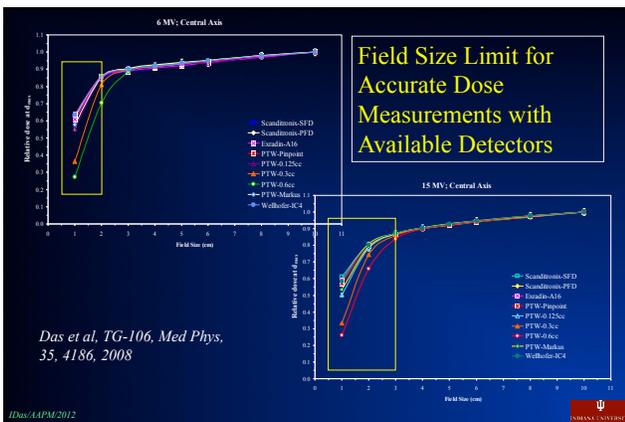
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## Relative Dosimetry

$$D_{w,Q_{msr}}^{f_{msr}} = M_{Q_{msr}}^{f_{msr}} N_{DW,Q_0} k_{Q,Q_0} k_{Q_{msr},Q}^{f_{msr},f_{ref}}$$

$$\Omega_{Q_{clin},Q_{msr}}^{f_{clin},f_{msr}} = \frac{M_{Q_{clin}}^{f_{clin}}}{M_{Q_{msr}}^{f_{msr}}} \left[ \frac{(D_{w,Q_{clin}}^{f_{clin}})/(M_{Q_{clin}}^{f_{clin}})}{(D_{w,Q_{msr}}^{f_{msr}})/(M_{Q_{msr}}^{f_{msr}})} \right] = \frac{M_{Q_{clin}}^{f_{clin}}}{M_{Q_{msr}}^{f_{msr}}} k_{Q_{clin},Q_{msr}}^{f_{clin},f_{msr}}$$

$$k_{Q_{clin},Q_{msr}}^{f_{clin},f_{msr}} = \frac{(D_{w,Q_{clin}}^{f_{clin}})/(M_{Q_{clin}}^{f_{clin}})}{(D_{w,Q_{msr}}^{f_{msr}})/(M_{Q_{msr}}^{f_{msr}})} = \frac{(Output)_{rel}}{(Reading)_{rel}}$$

$$k_{Q_{clin},Q_{msr}}^{f_{clin},f_{msr}} = \frac{(S_{w,air})_{f_{clin}} \cdot P_{f_{clin}}}{(S_{w,air})_{f_{msr}} \cdot P_{f_{msr}}}$$

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## New Data on Correction Factor

Table III. Ratio of detector readings  $M_{clin}^{det}/M_{msr}^{det}$  and correction factors  $k_{clin}^{det}/k_{msr}^{det}$  for the 5, 7.5, and 10 mm fields. Corresponding uncertainties at 68% level are shown in parentheses, indicating the uncertainty in the final one or two digits.

Detector	5 mm		7.5 mm		10 mm	
	$M_{clin}^{det}/M_{msr}^{det}$	$k_{clin}^{det}/k_{msr}^{det}$	$M_{clin}^{det}/M_{msr}^{det}$	$k_{clin}^{det}/k_{msr}^{det}$	$M_{clin}^{det}/M_{msr}^{det}$	$k_{clin}^{det}/k_{msr}^{det}$
A16	0.626 (15)	1.089 (3)	0.811 (10)	1.018 (3)	0.866 (6)	1.010 (3)
PinPoint	0.620 (17)	1.101 (3)	0.801 (7)	1.024 (3)	0.862 (5)	1.015 (3)
Diode 60008	0.726 (1)	0.943 (3)	0.873 (1)	0.949 (3)	0.912 (1)	0.964 (3)
Diode 60012	0.705 (1)	0.956 (3)	0.847 (2)	0.966 (3)	0.891 (1)	0.978 (3)
EDGE	0.726 (1)	0.948 (3)	0.864 (1)	0.955 (3)	0.906 (1)	0.966 (3)
Alanine	0.544 (8)	1.249 (8)	0.785 (12)	1.059 (4)	0.855 (13)	1.019 (5)
TLD	0.668 (4)	...	0.809 (6)	...	0.880 (8)	...
EBT films	0.659 (17)	...	0.811 (16)	...	0.853 (18)	...
Polymer gels <sup>a</sup>	0.702 (21)	...	0.872 (27)	...	0.929 (29)	...

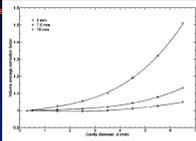


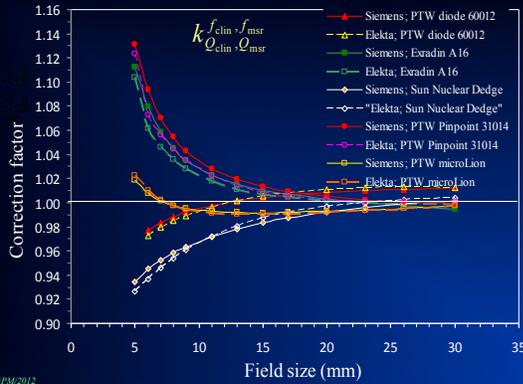
Fig. 5. Selected detector correction factors calculated using 40 percent gel dosimetry results (Ref. 9) and assuming cylindrical detectors of 2.5 mm length and varying active diameter for the 5, 7.5, and 10 mm field conditions for beam 6 MV x-rays SRS. The uncertainty of the presented results is of the order of 4% and the solid lines correspond to weighted least square regression fit results. In the absence of all other measurement perturbations, these data correspond to  $k_{clin}^{det}/k_{msr}^{det}$  values.

Table IV. Output factors  $k_{clin}^{det}/k_{msr}^{det}$  for the 5, 7.5, 10, and 15 mm fields. After correction factors are applied at all field sizes except 15 mm, as shown in Table III. Corresponding uncertainties at 68% level are shown in parentheses indicating the uncertainty in the final one or two digits.

Detector	5 mm	7.5 mm	10 mm	15 mm
A16	0.602 (17)	0.823 (10)	0.874 (7)	0.979 (3)
PinPoint	0.601 (18)	0.820 (8)	0.871 (5)	0.979 (3)
Diode 60008	0.698 (2)	0.829 (3)	0.879 (3)	0.980 (1)
Diode 60012	0.679 (2)	0.818 (3)	0.872 (3)	0.980 (1)
EDGE	0.699 (2)	0.823 (3)	0.873 (3)	0.980 (1)
Alanine	0.679 (11)	0.813 (13)	0.872 (13)	0.980 (1)
TLD	0.668 (4)	0.809 (6)	0.880 (8)	0.941 (7)
EBT films	0.659 (17)	0.811 (16)	0.853 (18)	0.955 (20)
Polymer gels <sup>a</sup>	0.702 (21)	0.872 (27)	0.929 (29)	...
Weighted mean	0.681 (1)	0.828 (1)	0.873 (1)	0.984 (1)

Pantelis et al. Med Phys. 37(6), 2369-2378, 2010

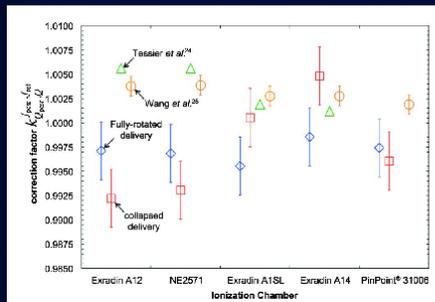
IBase/AAPM2012



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## New Data on Correction Factor

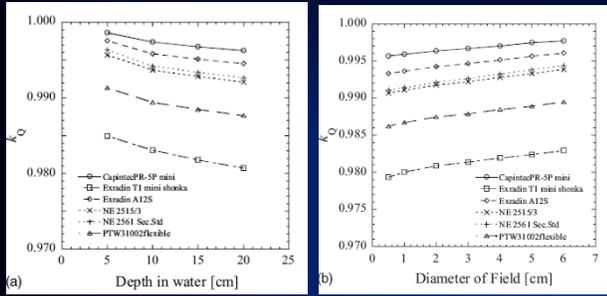


Chung et al. Med Phys. 37(6), 2404-2413, 2010

IBase/AAPM2012



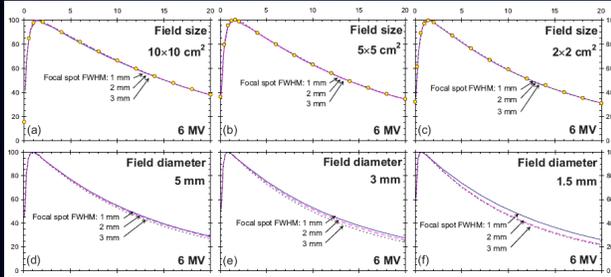
### $k_Q$ is not Constant in Small Field



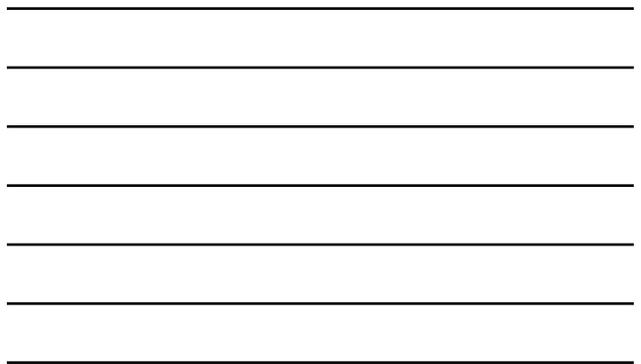
Kawachi et al, Med Phys, 35, 4591-4598, 2008



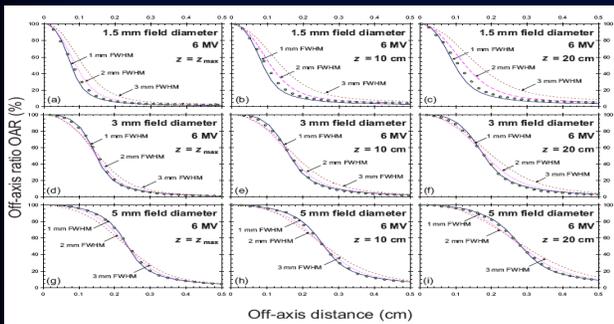
### Depth Dose & Source Size



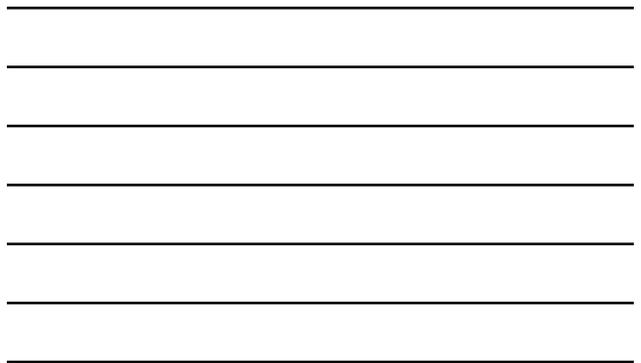
Sham et al, Med Phys, 35, 3317-3330, 2008



### Profile & Source Size

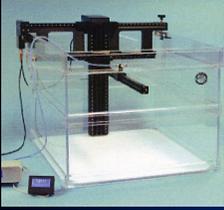


Sham et al, Med Phys, 35, 3317-3330, 2008



### Comparison of Large Tank and Small SRS Cylinder Tank for SRS, TMR & Profiles

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Moving Tank System for TMR

ARM Inc., Port Saint Lucie, FL 34983

IDash/AAPM/2012 PSI  
SUNSHINE UNIVERSITY

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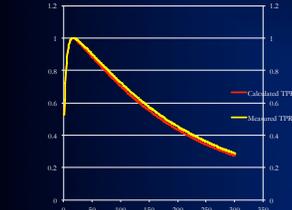
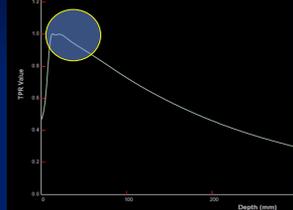
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### Direct TMR Data Acquisition

No SSD to SAD calculations required, No cubic spline fit of a limited number of fixed data points needed

Calculated TPR ~2% less at depth      Cubic spline fit of 12 data points

IDash/AAPM/2012 Nikesch et al, CyberKnife Center, Palm Beach, FL  
PSI  
SUNSHINE UNIVERSITY

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### 3D Scanner (Sun Nuclear)

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- ❖ Setup Subjectivity
  - ⌘ Automatic leveling, water surface detection and beam center detection
  - ⌘ No tank shifts
- ❖ Detector orientation/resolution
  - ⌘ 3D Scanner design always using the short dimension of chamber to scan
- ❖ Time
  - ⌘ Setup is faster and more accurate
  - ⌘ No tank shifts
  - ⌘ Smaller tank fills and drains faster

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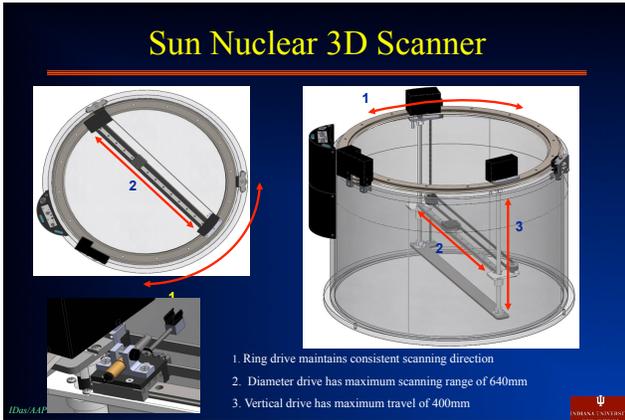
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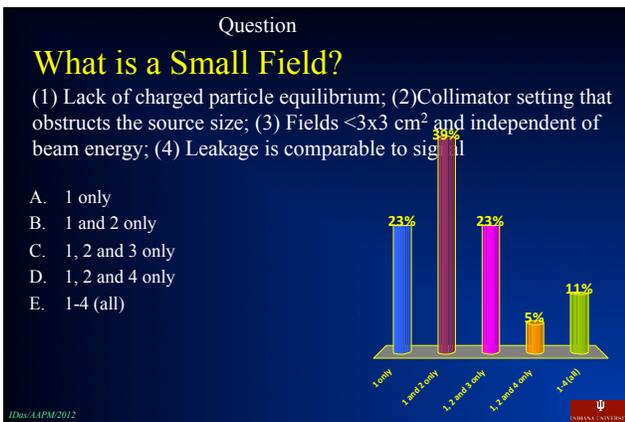
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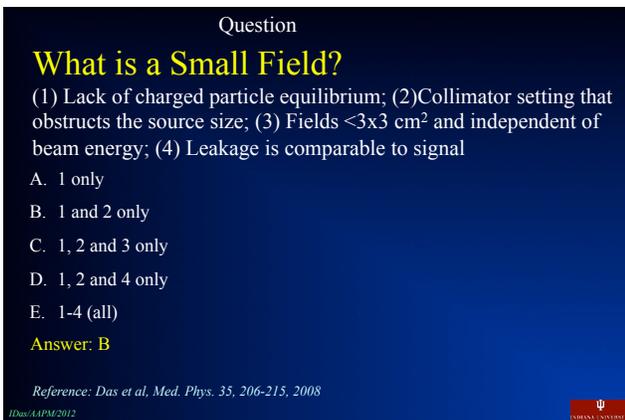
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Question

What is  $k_{Q_{clin}^{f_{msr}}}$  ?

- 17% A. Defined in TG-51 for chamber correction factor
- 46% B. Defined in noncompliant TG-51 dosimetry for correcting reading to actual dose
- 33% C. Conversion factor from  $K_Q$  to Dose
- 0% D. Used in dynamic and Arc therapy dose calculation
- 4% E. Used primarily in Tomotherapy dosimetry

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Question

What is  $k_{Q_{clin}^{f_{msr}}}$  ?

- A. Defined in TG-51 for chamber correction factor
- B. Defined in noncompliant TG-51 dosimetry for correcting reading to actual dose
- C. Conversion factor from  $K_Q$  to Dose
- D. Used in dynamic and Arc therapy dose calculation
- E. Used primarily in Tomotherapy dosimetry

Answer: B

Reference: Alfonso, et al. Med Phys 35, 5179-5186 (2008)

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## Conclusions

- ❖ Golden Data should be taken as a reference only
- ❖ Understand time and amount of data to be taken
- ❖ View each parameters properly, double check by another individual
- ❖ Use proper detector for each type of data collection
- ❖ Set optimum speed for scanning, do not rush

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## -Conclusions

- ❖ Understand the limits and measuring condition
- ❖ Question every unusual data set
- ❖ Do not smooth data too much
- ❖ Write report for future reference
- ❖ Future technology & resources could help commissioning simpler

Das/AAPM2012



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