

Optimizing dose and image quality in fluoroscopy

A. Kyle Jones, Ph.D.

Assistant Professor

MD Anderson Cancer Center

Disclosure

- I am co-owner of Fluoroscopic Safety, LLC, a company that markets educational programs on the safe use of fluoroscopy

Educational objectives

- Identify targets for dose and image quality optimization in fluoroscopy
- Describe strategies for exploiting these targets
- Realize that dose optimization must be considered in the broader context of procedural goals

Optimize, not reduce

- We should always be speaking in terms of *optimizing* clinical procedures that are *justified*
- The use of “dose reduction” implies efforts to reduce dose without consideration of image quality
- Especially in fluoroscopy, attempting to reduce dose with a blind eye towards image quality can actually *increase* dose in the end

Targets for optimization

- Equipment configuration/calibration
- Practice/use of fluoroscopy
- Technology

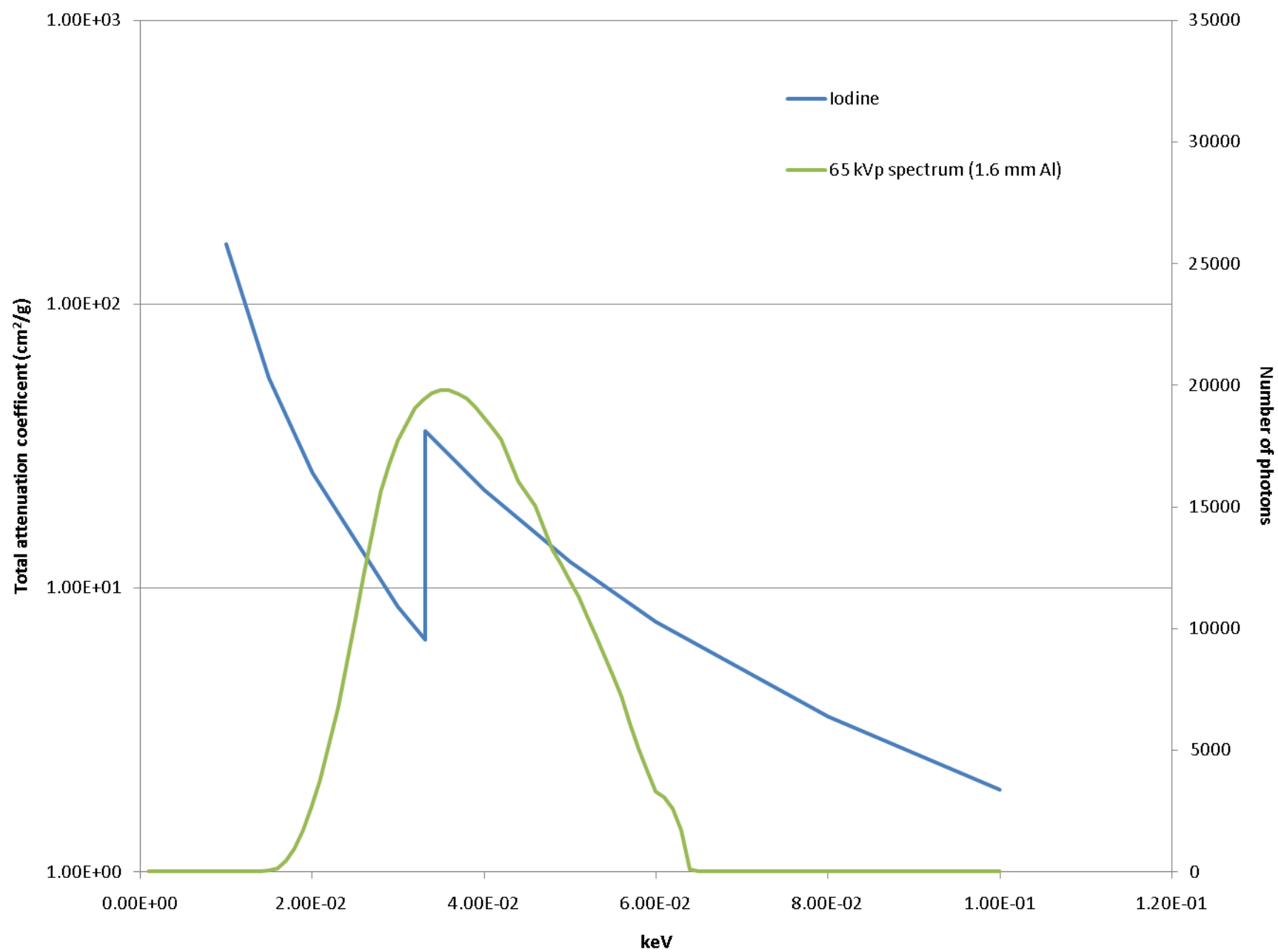
EQUIPMENT CONFIGURATION AND CALIBRATION

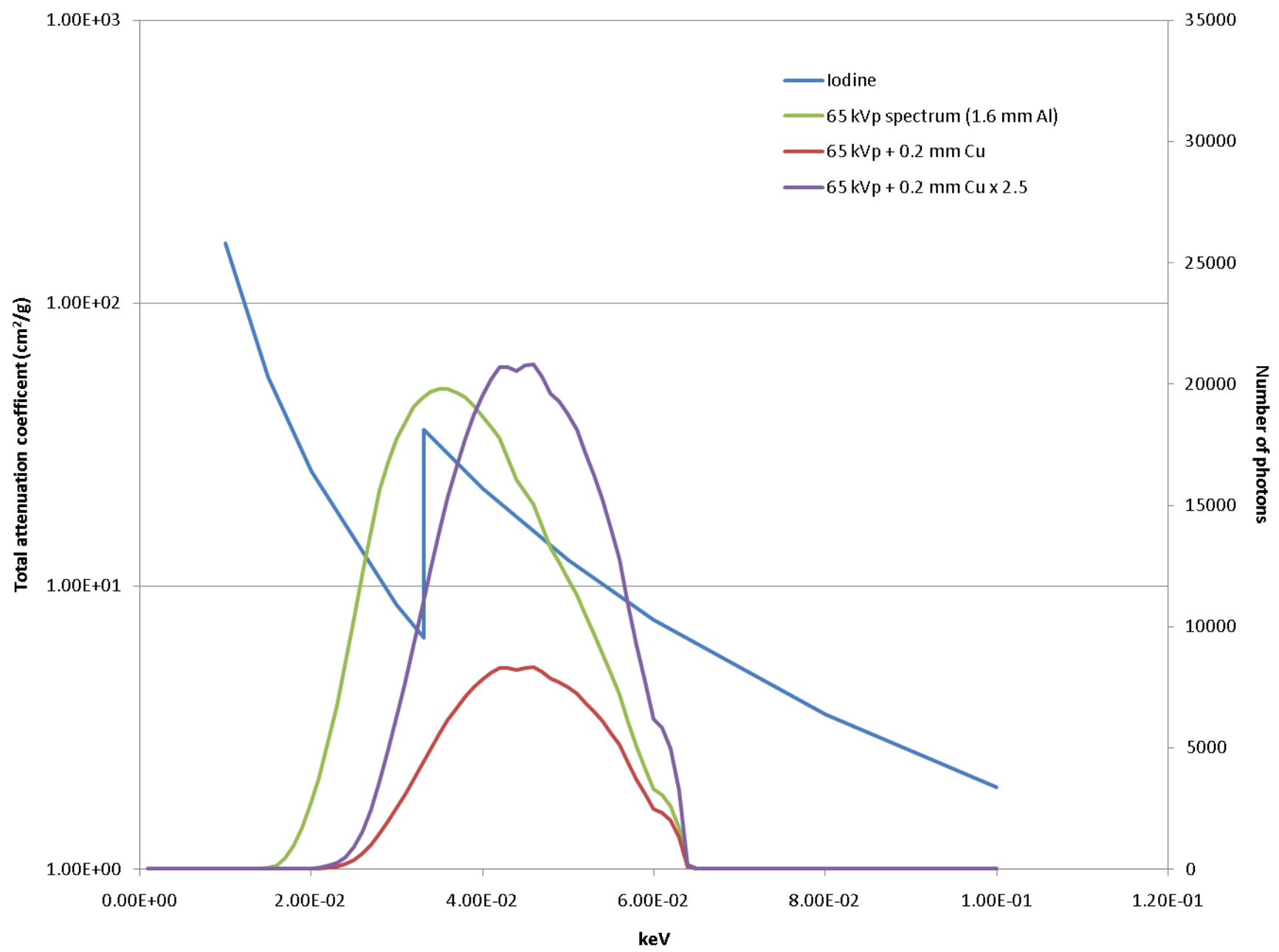
Easy targets for optimization

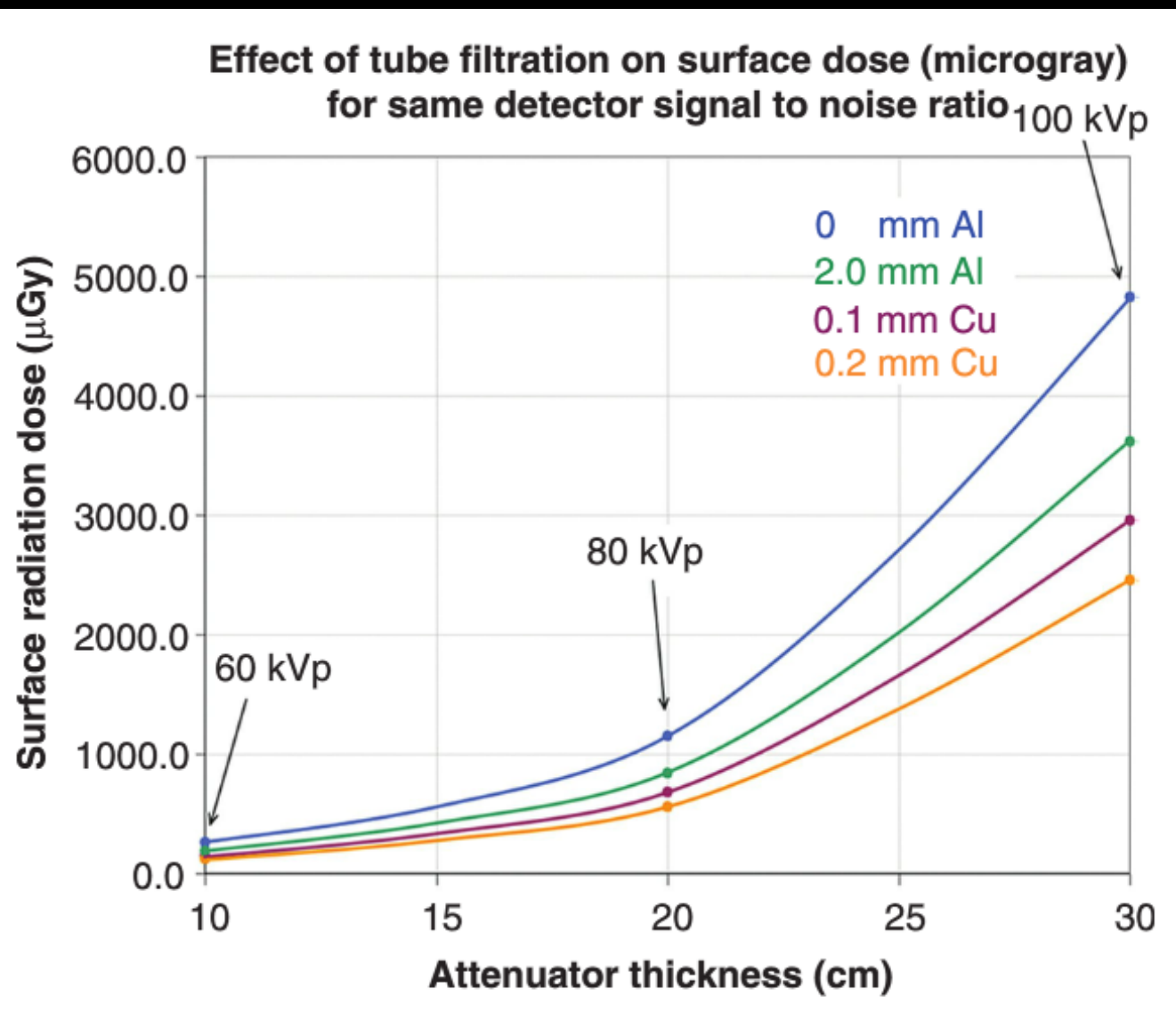
- Minimum filtration
- Positioning without radiation configured to be enabled by default
- Configure desired organ program to be loaded by default
- Use of variable frame rate
- Mask averaging
- Choose the right equipment
- Use pulsed fluoroscopy and configure it to use the Aufrichtig scale



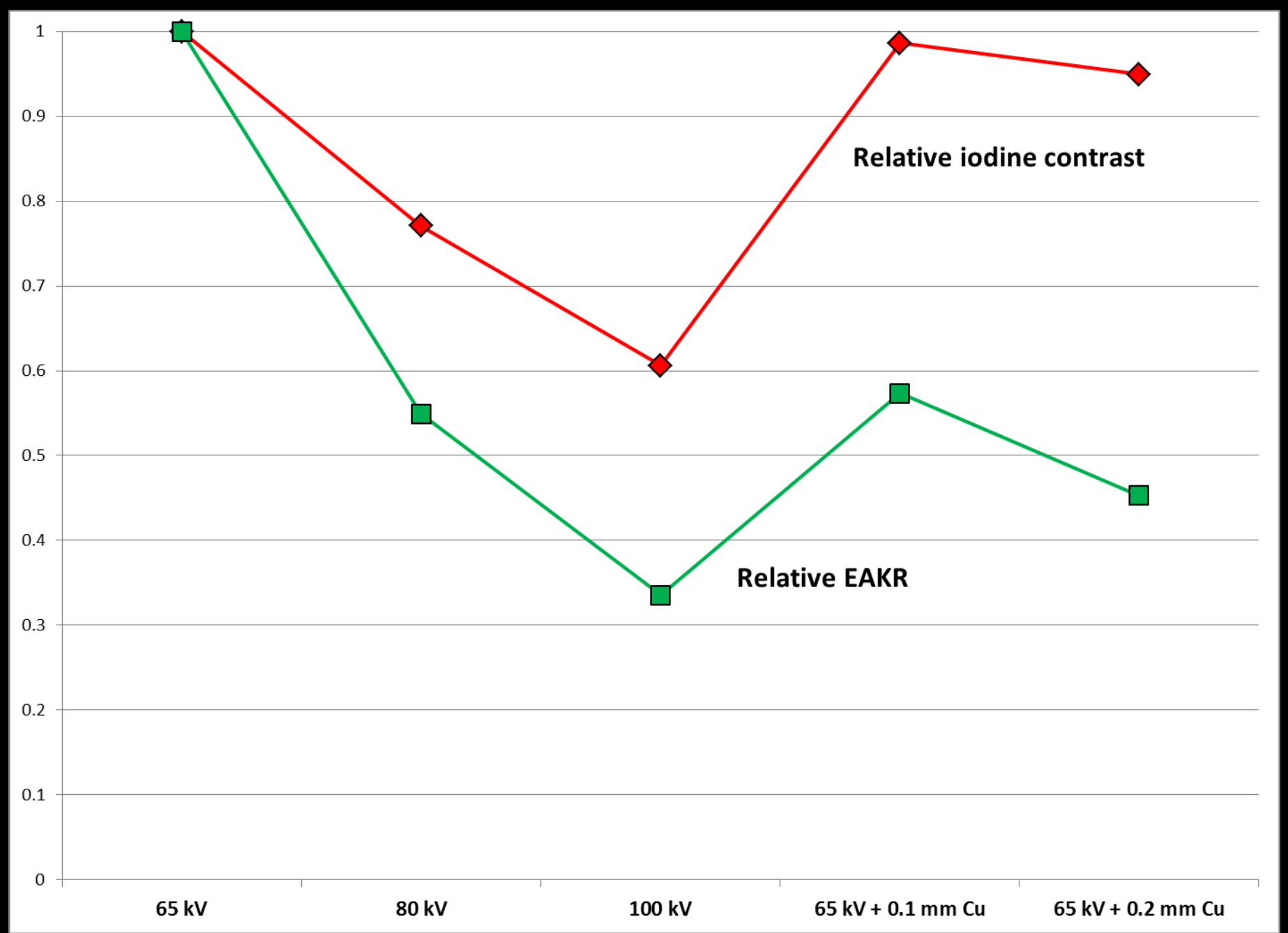
Scott Adams, www.dilbert.com





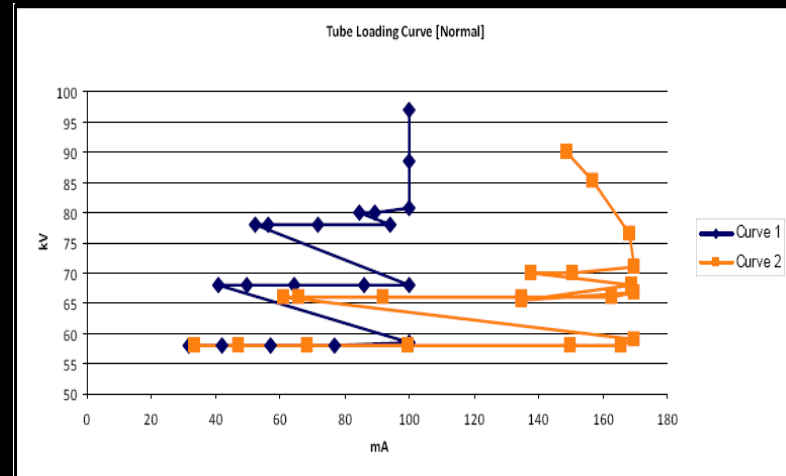


Bushberg et al. The Essential Physics of Medical Imaging, 3rd ed.



Contrast and beam quality

- Iodine contrast is strongly affected by beam quality
- The addition of filtration allows the use of low kV while maintaining dose at an acceptable level
 - Traditional, Program-Switched
- Sacrifices may need to be made to maintain kV at desired level
 - E.g., focal spot



Ishida E et al. Image quality improvement and patient dose reduction by optimization of x-ray spectrum. Jpn Radiol Technol **55**:582-587 (1999).

Balancing filtration

- Impact in FLU vs. ACQ
- Must consider impact on related organ program parameters
 - Pulse width
 - Focal spot size
- Configuration options
 - “Traditional” method
 - Static filter
 - “Program-switched” method
 - Filtration linked to organ program (e.g., Philips)
 - Seissl method (e.g., Siemens)

FLUORO	(2) Cardiac Cath		
	FL Card Low (21,4)	FL Card Avg (21,5)	FL Card Lg (21,6)
EXPOSURE	FLUORO	FLUORO	FLUORO
kV	81	70	70
Pulse Width	8	8	8
kV ms	90	90	90
kV dose	109	109	off
EP Reduction	2	2	
<i>Cu mm Min</i>	0.6	0.2	0.1
<i>Cu mm Max</i>	0.9	0.6	0.3
<i>Dose</i>	36	45	55
<i>High Contrast (20R)</i>	Normal	Normal	High
<i>kV Warning</i>	109	109	off
<i>Focus</i>	s	s	s

Exam Set and Program Editor

Nephro - Prone

CARE Single (Single)

FL Angio 7.5 (7.5 p/s)

<< Standard

Display Fluoro

Display Roadmap

FL Angio 7.5 (7.5 p/s)

FL Angio 15 (15 p/s)

DynaCT Body

Biliary

Venogr.

Nephro - Prone

CARE Single (Single)

Native (1 f/s)

Native 2 (2 f/s)

Sub 1 (1 f/s)

FL Angio 10 (10 p/s)

FL Angio 7.5 (7.5 p/s)

FL Angio 15 (15 p/s)

Nephro - Supine

Native

CARE Single (Single)

Native (1 f/s)

Native 2 (2 f/s)

Sub 1 (1 f/s)

FL Angio 10 (10 p/s)

FL Angio 7.5 (7.5 p/s)

FL Angio 15 (15 p/s)

CO 2/GAD

General DSA

Pelvis

Pulmonary

IVC

Program FL Angio 7.5

Mode FLUORO

KV	70 KV
Pulsewidth	10.0 ms
KV ms	96 KV
Focus	
Dose	45 nGy/p
KV Dose	109 KV
EP Reduction	2.0 EP
Skindose Profile	Normal Contrast
Min. CU-Filter	0.2 mm
Max. CU-Filter	0.9 mm
KV Warning Level	Off
I-Noise Reduction	Off
Edge Enhancement NAT	30 %
EE-Kernel	5
DDO	40 %
DDO-Kernel	137
Window Center	1800
Window Width	2500
Auto Window	<input checked="" type="checkbox"/>
Auto Window Setting	Normal(C=300,V
Auto-Window Center Correction	-150
Auto-Window Width Correction	1.0
Sigmoid Window	<input type="checkbox"/>
Gamma Correction	G06/C3
Gain Correction	0.0 EP
K-Factor	Auto7
EVE	Auto7

Activate Exam Set and Program

Remove from Exam Set

Save as New Exam Set...

Apply Parameters

Save as New Program

Save Parameters

Save and Apply

Login

Close

Help

13.03.21-08:41:...

1/1/1900

mGy A: 0

μGym² A:0.00

Nephro - Prone

DR Fixed

CARE Single

kV 70.0

mA 443.6

Measure Field

ms 10.7

Focus

Time

Cu mm 0.0

f/s Single

Dilatation 00:00

Σ A+B 000.0

FL Angio 7.5

kV 65.0

mA 168.5

ms 10.2

Cu mm 0.6

p/s 7.5 p/s

Heat Unit % 0 %

REF

AS

REF

SM

REF

Examination

PostProc

Quant

Filming

5 Worklist Item(s) received.

Exam Sets

1 HAE 2K	HEPATIC VFR Manual	CELIAC VFR Manual	SMA VFR Manual	GDA VFR Manual
	-----	Fluoro Angio 10 P/s	Fluoro Angio 7.5 P/s	Fluoro Angio 15 P/s
2 HAE HC	HEPATIC VFR Manual	CELIAC VFR Manual	SMA VFR Manual	GDA VFR Manual
	-----	Fluoro Angio 10 P/s	Fluoro Angio 7.5 P/s	Fluoro Angio 15 P/s
3 DynaCT	HEPATIC VFR Manual	5s-1k DR 1.5 °/F	8sDR 0.5 °/F	8s_DR_HQ 0.5 °/F
	-----	Fluoro Angio 10 P/s	Fluoro Angio 7.5 P/s	Fluoro Angio 15 P/s
4 Venogr.	Veno 1 1 F/s	Veno 2 2 F/s	DSA 3 3 F/s	Veno Single
	-----	Fluoro Angio 10 P/s	Fluoro Angio 7.5 P/s	Fluoro Angio 15 P/s
5 Biliary	PTC/Drain 1 F/s	PTC Sub 1 F/s	Native 2 2 F/s	Single Single
	-----	Fluoro Angio 10 P/s	Fluoro Angio 7.5 P/s	Fluoro Angio 15 P/s

Profile:

UNIVERSAL

Display
Roadmap

Delete

Save As New

Edit Prog...

Apply

Programs:

Acq

Show List...

SIEMENS

Close Editor

Help

Variable frame rate (VFR) imaging

- The concept of this technique is quite simple – by reducing the total number of ACQ frames, dose is reduced
- Frame rate reduction is triggered by
 - Time
 - Manually

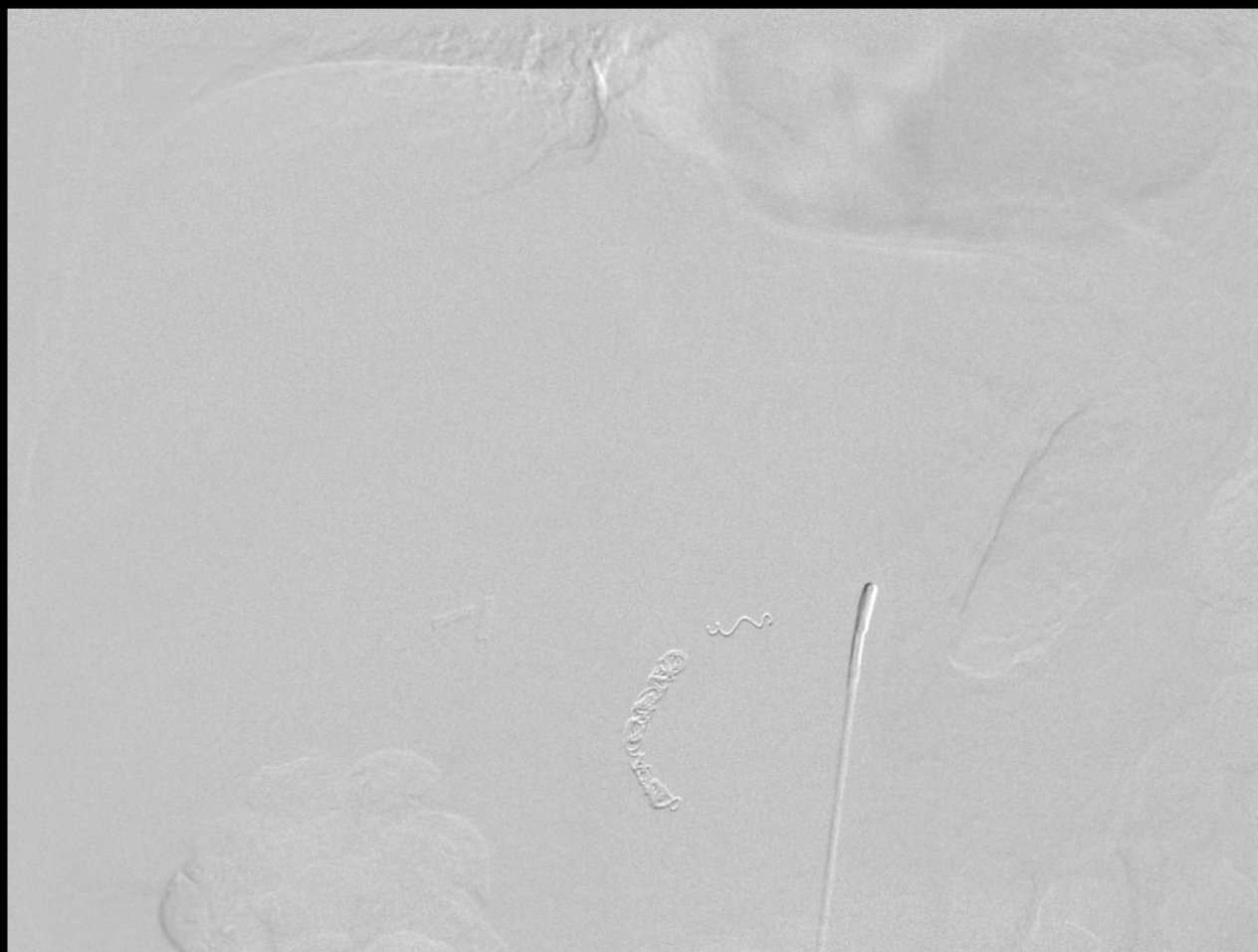
ACQUISITION TYPE

Scene Time: 40 s

Framerate Ctrl: VFR Manual

No Phases: 3

Phase	Framerate [f/s]
Phase 1	4 F/s
Phase 2	2 F/s
Phase 3	1 F/s

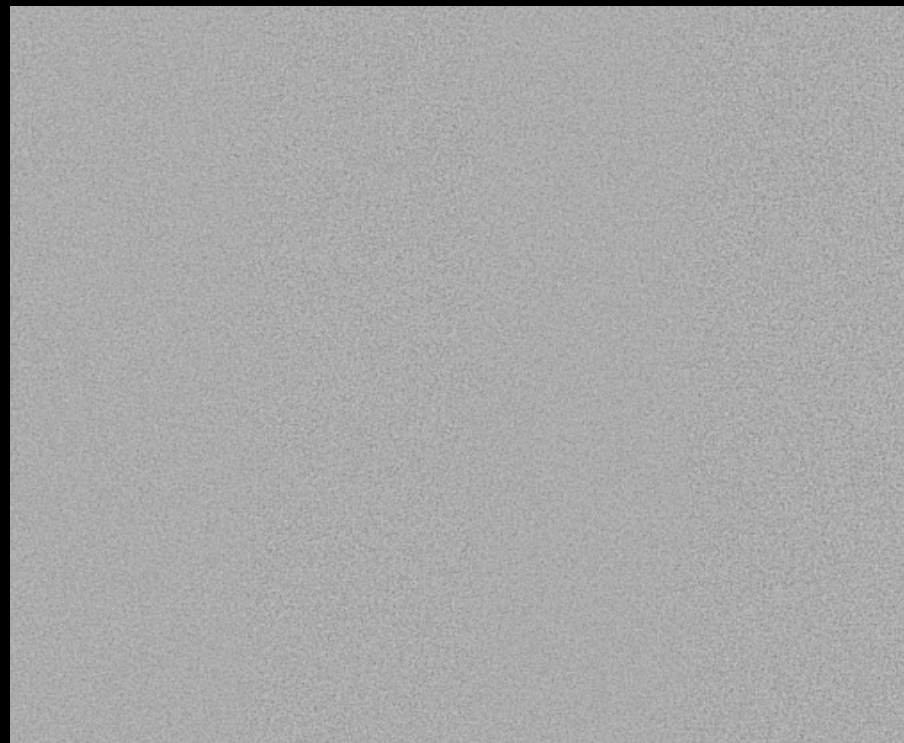


Mask averaging

- Reduction in image noise that can translate directly in to reduction in dose with virtually no impact on image quality
- Unfortunately, in my practice it is rarely used



3.6 uGy/fr



2.4 uGy/fr

Aufrichtig scale for pulsed fluoroscopy

- The human visual system has a finite integration time of approximately 200 ms
- Richard Aufrichtig studied this phenomenon and derived an empirical relationship relating the necessary dose per pulse to the pulse rate
- The use of lower pulse rates results in an LIH image of higher quality

$$\left(\frac{IAKRD_2}{IAKRD_1} \right) = \sqrt{\frac{PPS_1}{PPS_2}}$$

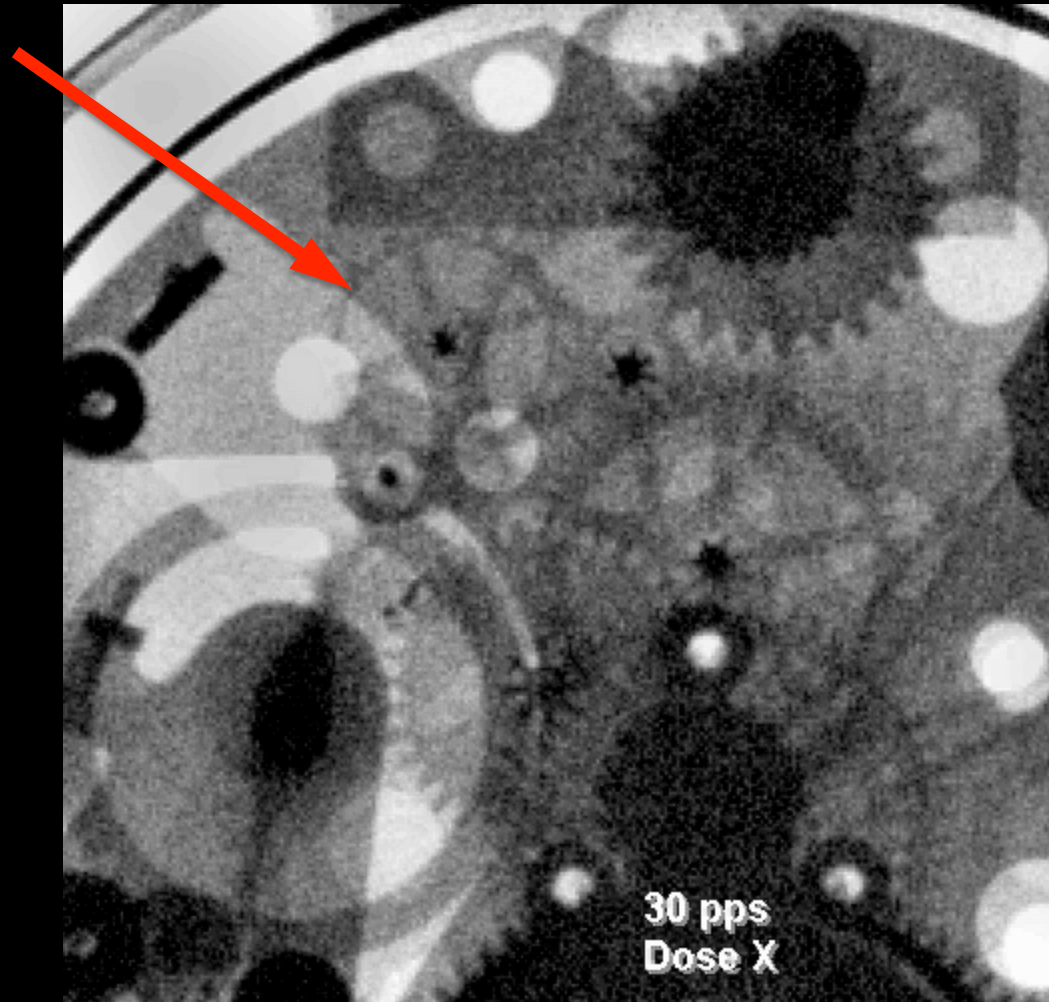


Illustration courtesy of Phil Rauch, M.S.

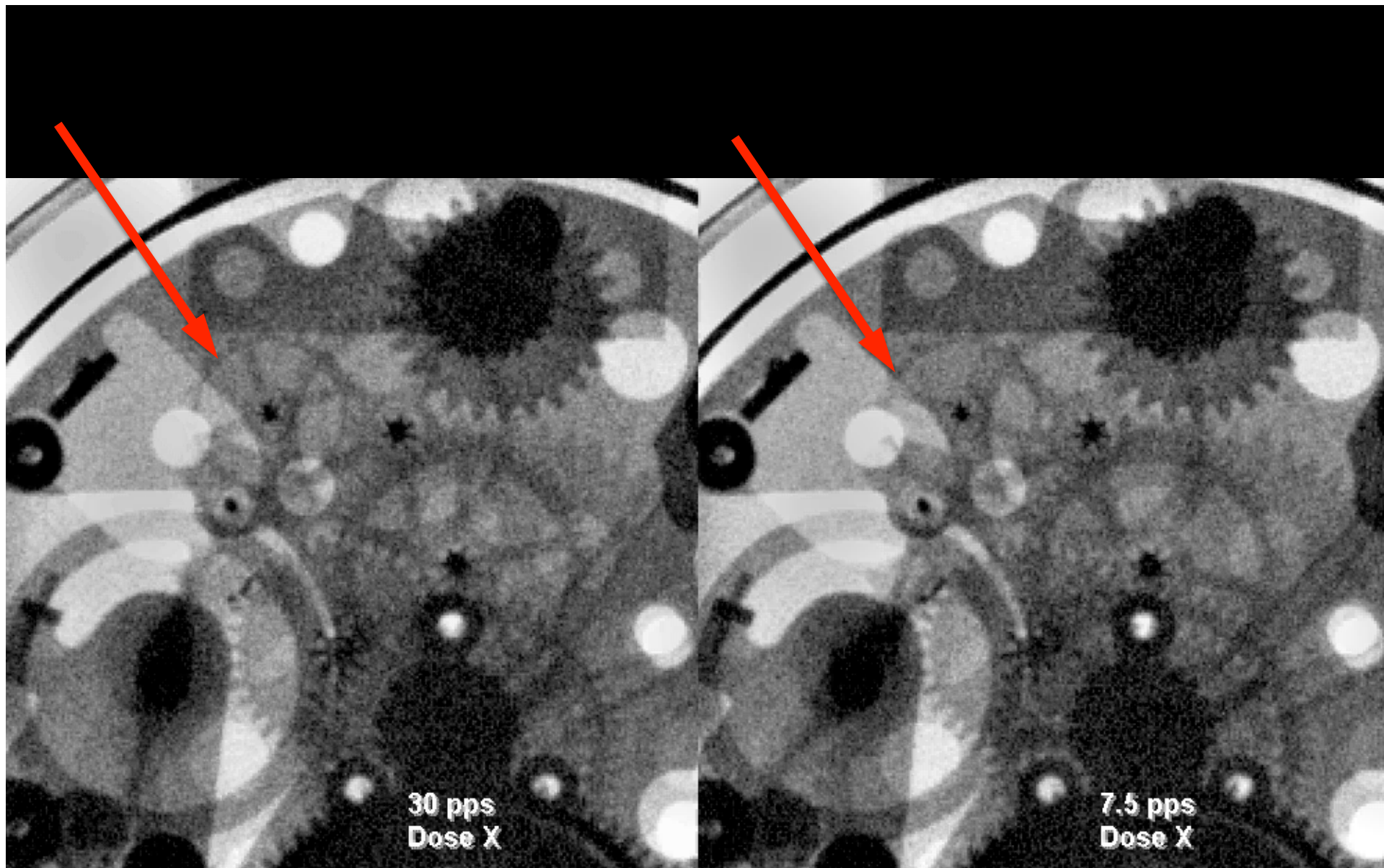


Illustration courtesy of Phil Rauch, M.S.

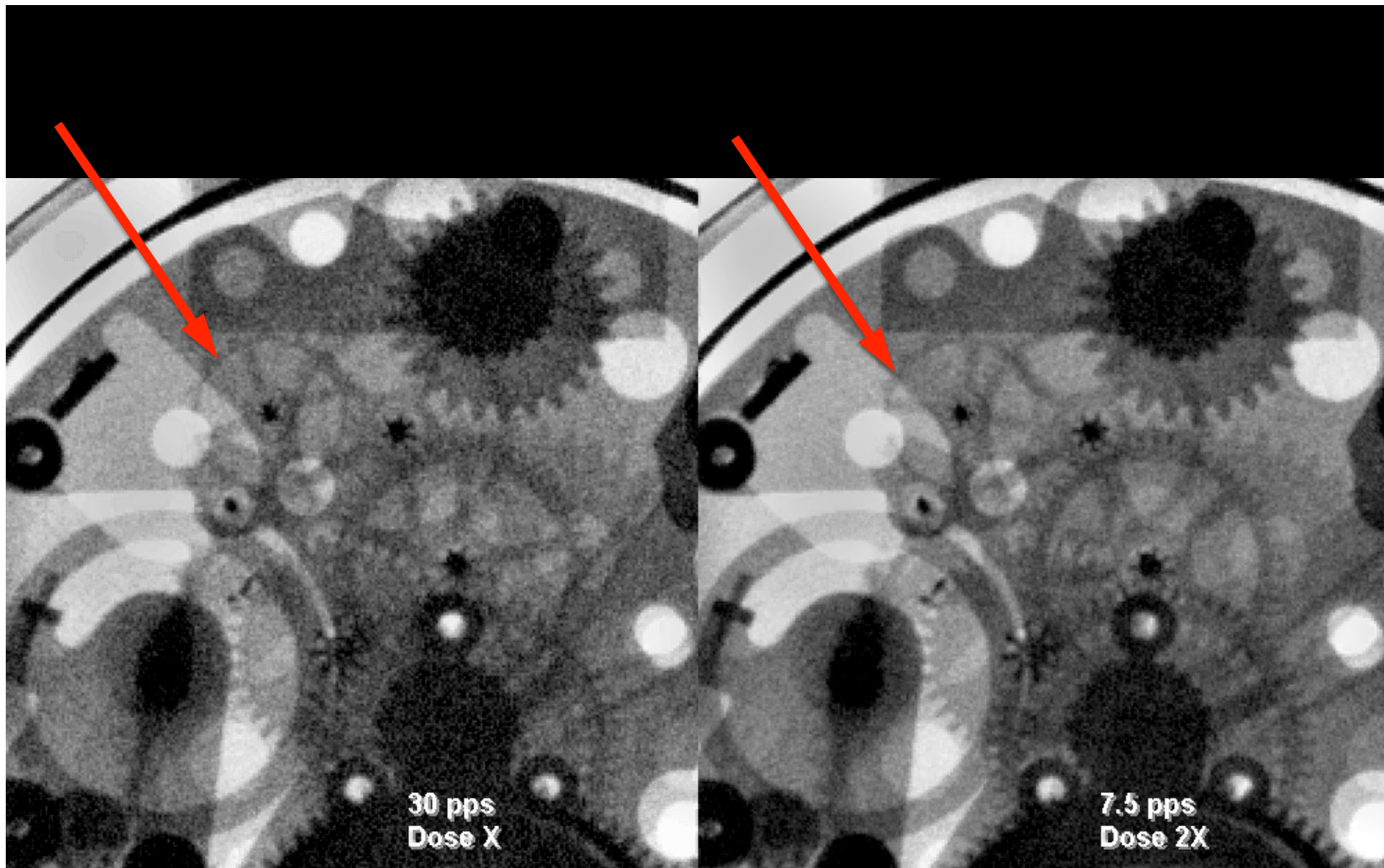


Illustration courtesy of Phil Rauch, M.S.

PHYSICIAN PRACTICE

Physician practice

- Physicians have plenty on their mind during a complex FGI
- We cannot expect them to also retain detailed knowledge of equipment settings and apply this knowledge during a case
- We should provide simple instructions and configure protocol defaults with this in mind
- *Speak their language*

Key aspects

- The Tetrad
- Notification levels and associated actions
- Dose audits

The “Tetrad”

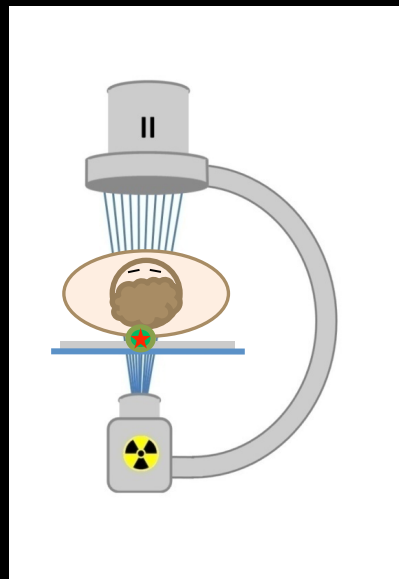
1. Raise the patient table to the highest comfortable working height.
2. Lower the image receptor as much as practicable.
3. Take one small step back or down the table away from the patient.
4. Collimate the X-ray field to the area of interest.

Order is important!

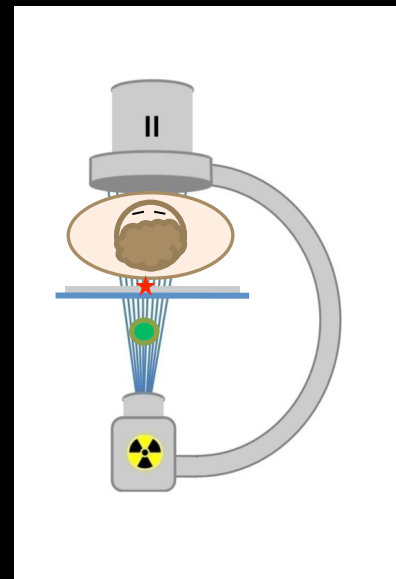
Procedural geometries

Using Good Practice is essential during fluoroscopic procedures, including maintaining the patient as far from the X-ray source as practical. This is common in vascular and interventional procedures, and results in a *non-isocentric* geometry.

The dose rate at the skin surface will be lower in a non-isocentric geometry than in an isocentric geometry.



Isocentric geometry



Non-isocentric geometry

TABLE 4.7—*Suggested values for first and subsequent notifications and the SRDL.*

Dose Metric	First Notification	Subsequent Notifications (increments)	SRDL
$D_{\text{skin,max}}$	2 Gy	0.5 Gy	3 Gy
$K_{\text{a,r}}$	3 Gy	1 Gy	5 Gy ^a
P_{KA}	300 Gy cm ² ^b	100 Gy cm ² ^b	500 Gy cm ² ^b
Fluoroscopy time	30 min	15 min	60 min

^aSee additional discussion concerning the value 5 Gy in Section 4.3.4.2.

^bAssuming a 100 cm² field at the patient's skin. For other field sizes, the P_{KA} values should be adjusted proportionally to the actual procedural field size (*e.g.*, for a field size of 50 cm², the SRDL value for P_{KA} would be 250 Gy cm²).

NCRP 168

Notification levels by lab/fluoroscope type

Differences in notification levels reflect differences, technical and geometric, in how fluoroscopically-guided procedures are performed. A number of factors influence the ratio of peak skin dose (PSD) to RAK: procedural geometry, backscatter, and attenuation by the patient support and pad.

Each notification level should involve a procedural pause and communication of the radiation dose status to the operator.

Lab/fluoroscope type	Typical ratio of PSD to RAK (PSD/RAK)
Vascular/interventional radiology	1.0
Cardiac catheterization	1.3-1.4
Interventional neuroradiology	1.0-1.3
Mobile C-arm	1.0-1.5

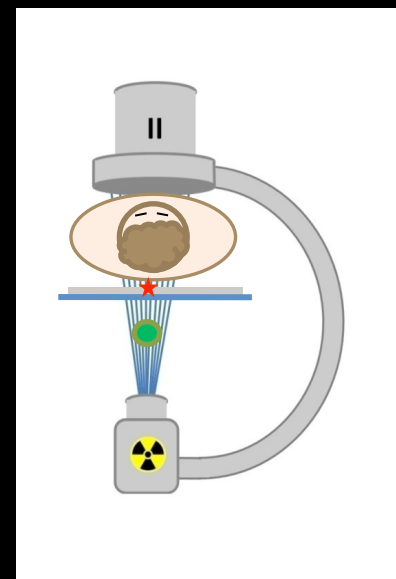
Establishing a Patient Safety Program in Fluoroscopy, Fluoroscopic Safety, LLC.

Example – Vascular/Interventional Radiology

RAK notification level (mGy)	Corresponding PSD (mGy)	Suggested action
2,500	2,500	Verify Good Practice is being used.
5,000	5,000	Substantial radiation dose level. Flag patient for f/u. Measure and record table height.
7,500	7,500	Verify Good Practice. Re-evaluate risk/benefit pace of procedure, entering range of potential skin injury.
10,000	10,000	Verify Good Practice. Re-evaluate risk/benefit pace of procedure. Skin injury more likely.

**And every 1,000 mGy above 10,000 mGy. These notification levels are for illustration purposes only and the numbers are approximate.*

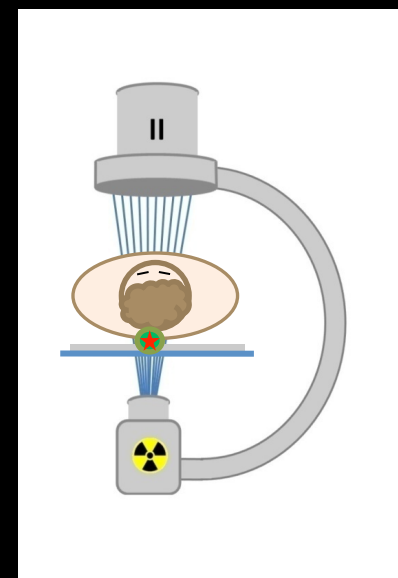
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The use of a non-isocentric geometry means that PSD is often similar to RAK for VIR procedures.

Example – Interventional cardiology

RAK notification level (mGy)	Corresponding PSD (mGy)	Suggested action
1,800	2,500	Verify Good Practice is being used.
3,600	5,000	Substantial radiation dose level. Flag patient for f/u. Measure and record table height. Consider rotating C-arm.
5,400	7,500	Verify Good Practice. Re-evaluate risk/benefit pace of procedure, entering range of potential skin injury.
7,200	10,000	Verify Good Practice. Re-evaluate risk/benefit pace of procedure. Skin injury more likely. Consider rotating C-arm.



The use of an isocentric geometry means that PSD is often greater than RAK for interventional cardiology procedures.

**And every 700 mGy above 7,200 mGy. These notification levels are for illustration purposes only and the numbers are approximate.*

Establishing a Patient Safety Program in Fluoroscopy, Fluoroscopic Safety, LLC.

Dose audits

- The simple act of beginning to record dose metrics will often on its own trigger practice changes
- Stratification of the data can identify specific targets for improvement
- Compare to normative data sets
 - E.g., RAD-IR study

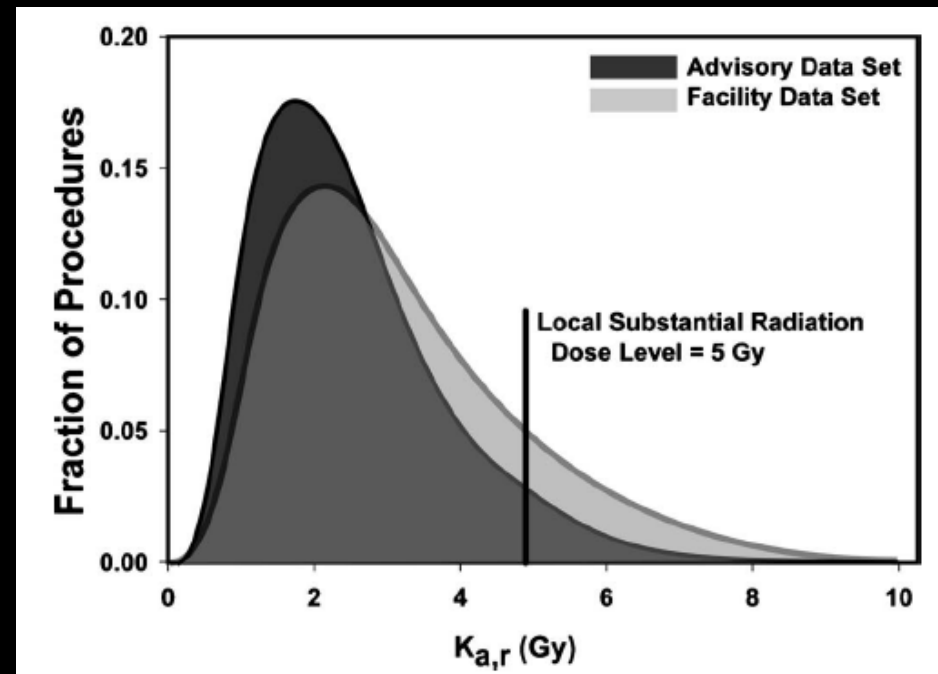
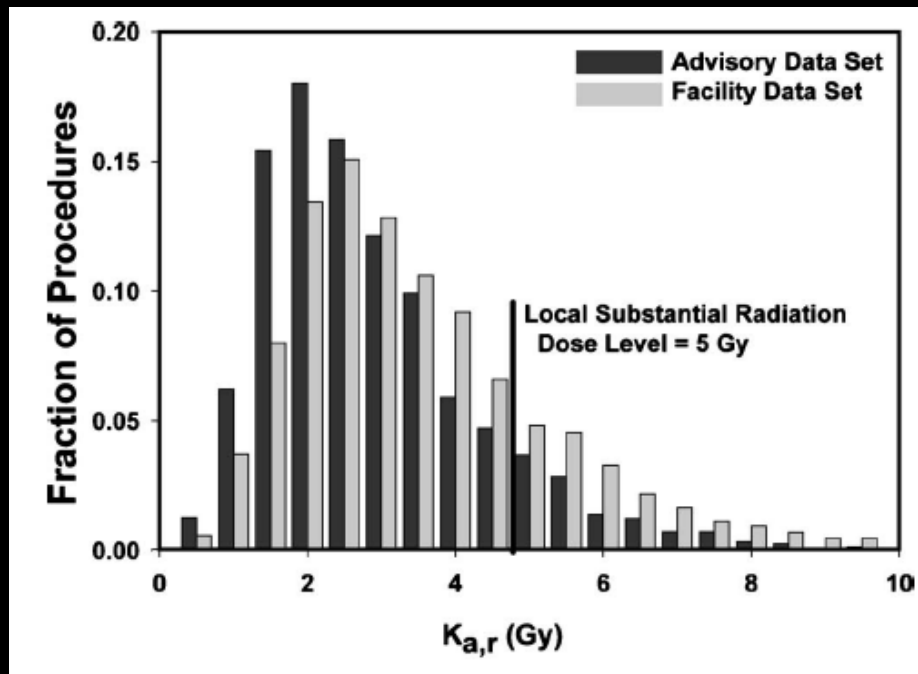
Recommendation 19

Facilities *shall* have a process to review radiation doses for patients undergoing FGI procedures.

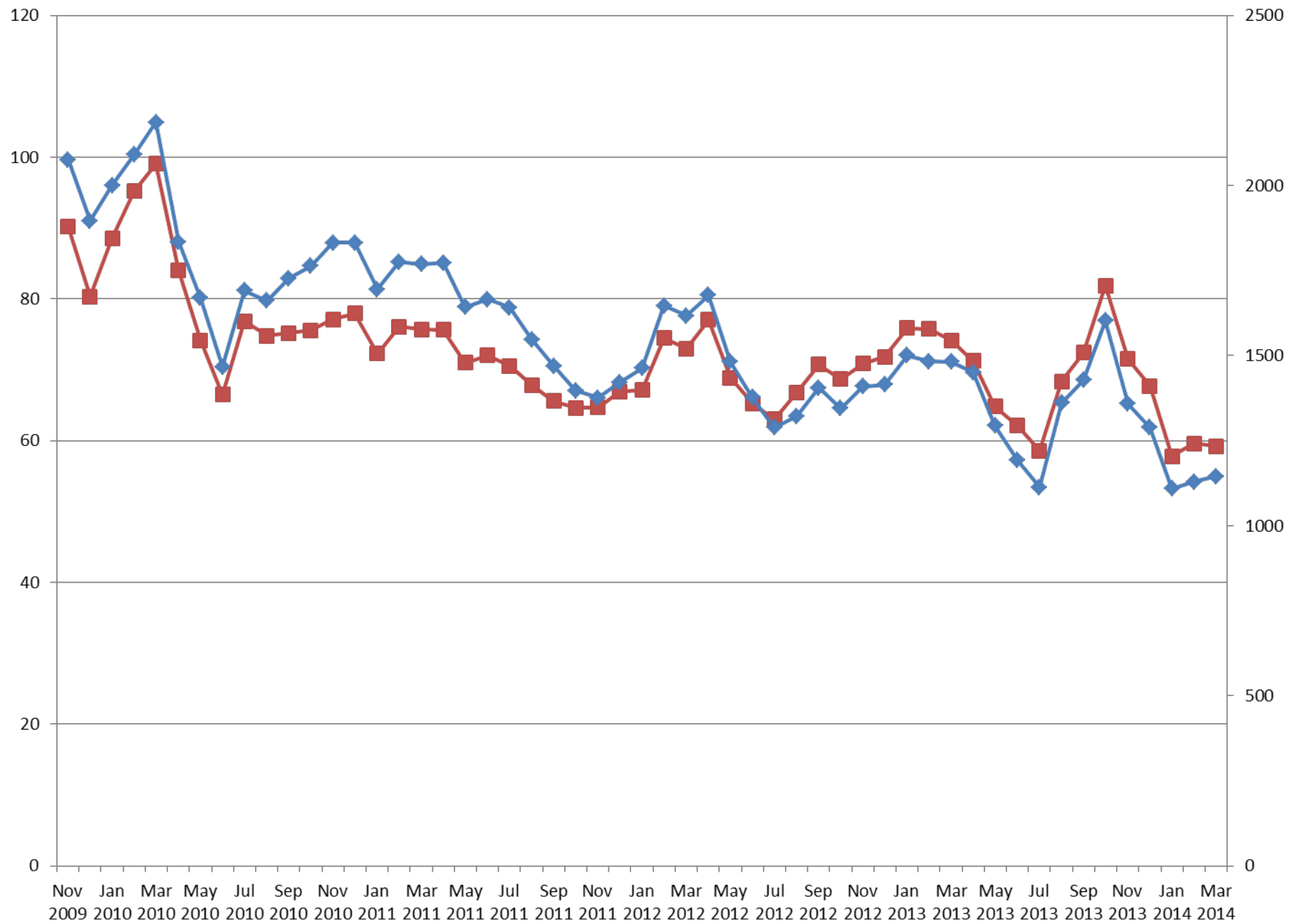
Advisory data based on measured dosimetric quantities (in particular P_{KA} or $K_{a,r}$ to manage overall performance, and $K_{a,r}$ to manage deterministic effects) *should* be used for quality assurance purposes.

NCRP 168

Radiation dose audits



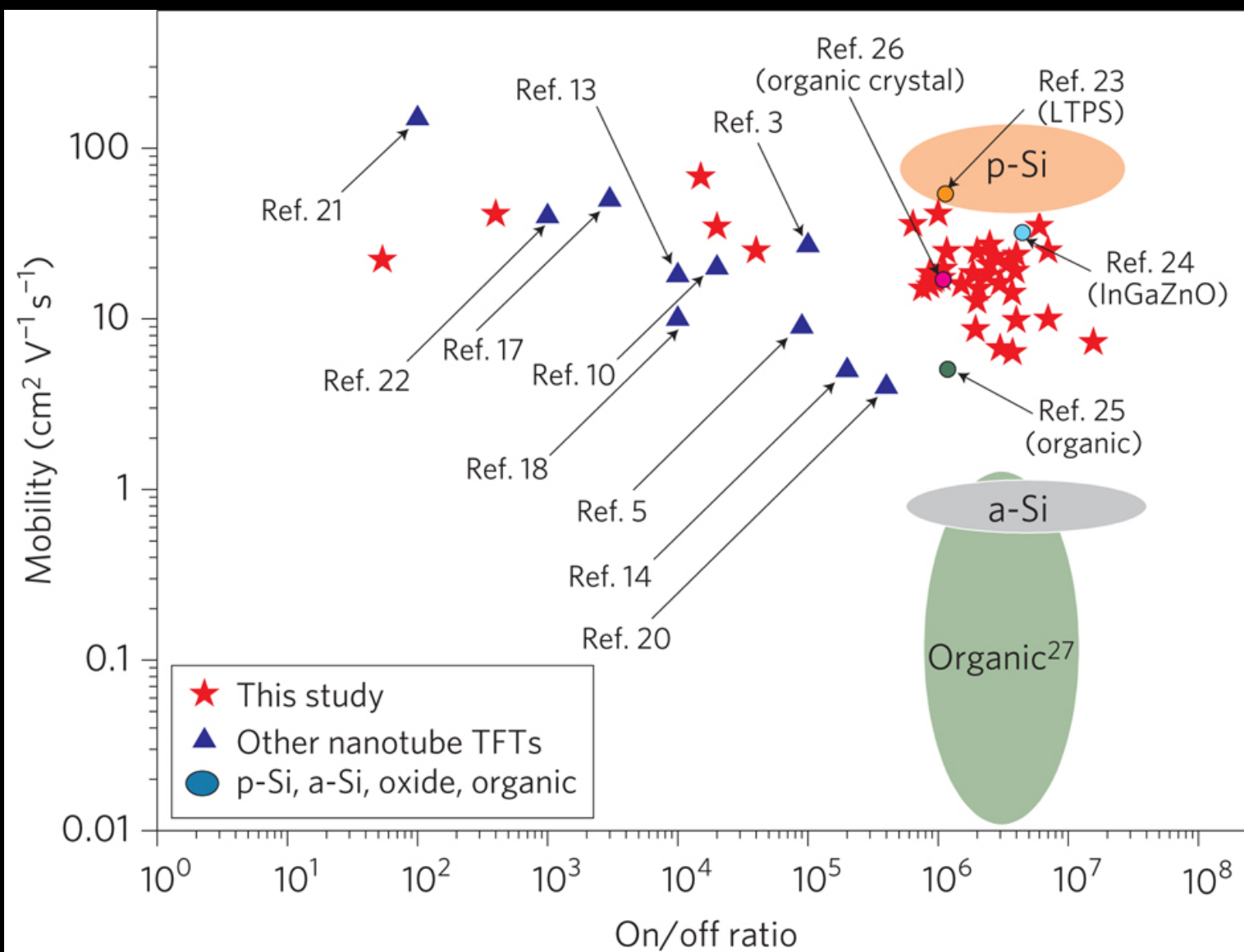
Balter S et al. Patient radiation dose audits for fluoroscopically guided interventional procedures. Med Phys 38(3):1611-18, 2011.



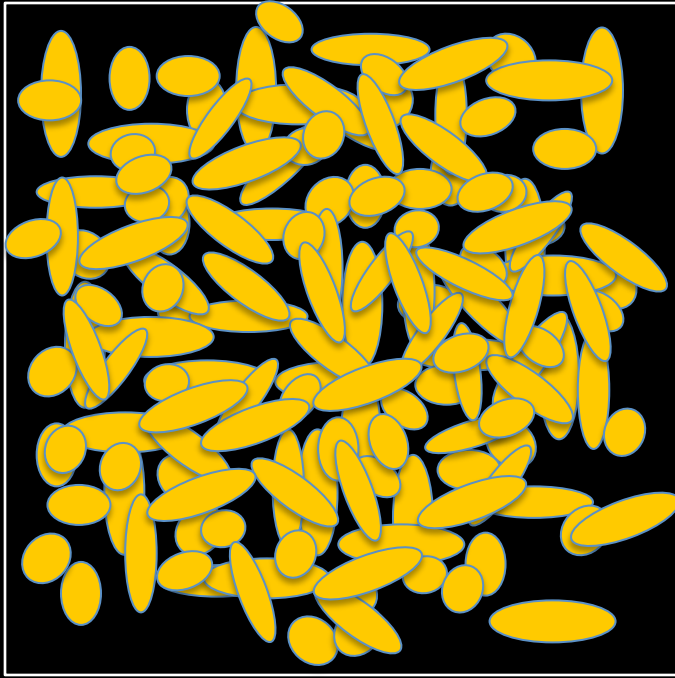
TECHNOLOGY

Technological advances

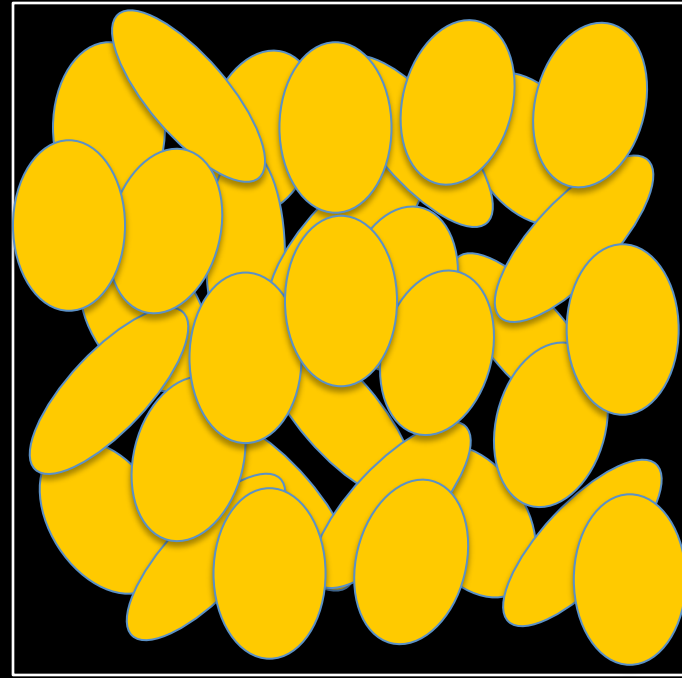
- Detector technology
 - Higher acquisition bit depth
 - Crystalline silicon
 - Electron mobility
 - Active pixel sensors
- X-ray tube technology
 - Flat emitter



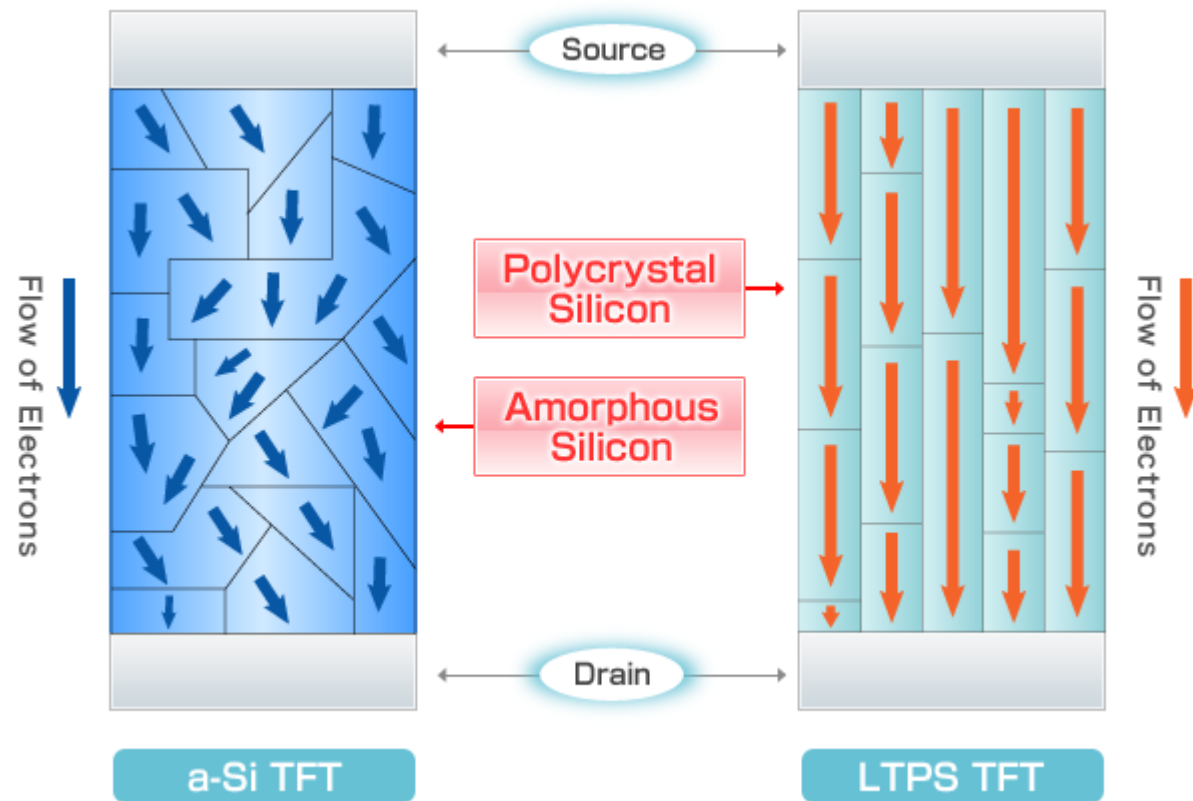
Sun D et al. Flexible high-performance carbon nanotube integrated circuits. Nat Nanotechnol 6:156-161 (2011).



a-Si



poly-Si



Backplane	a-Si:H	poly-Si	mono-Si
Monolithic array size	Large	Medium	Small
e ⁻ mobility	1 cm ² /V-s	100 cm ² /V-s	1000 cm ² /V-s
TFT mask steps	4-5	5-11	5-11
Leakage current	Low	Higher	Higher
Radiation hardness	Excellent	Good/Fair	Poor
Large scale uniformity	Good	Fair	Poor
Cost/yield	Low/High	High/Low	High/Low

Further reading

- Miller et al., Quality improvement guidelines for recording patient radiation dose in the medical record. *J Vasc Interv Radiol* **15**:423–429, 2004.
 - SIR Standards of Practice Committee
- Miller DL, Balter S, Noonan PT, Georgia JD, Minimizing radiation-induced skin injury in interventional radiology procedures. *Radiology* **225**:329–336, 2002 .
- Stecker et al., Guidelines for patient radiation dose management. *J Vasc Interv Radiol* **20**:S263–S273, 2009.
 - SIR Safety and Health Committee
 - Discharge/consenting examples
- Archer BR and Wagner LK, Protecting patients by training physicians in fluoroscopic radiation management. *J Appl Clin Med Phys* **1**:32-37, 2000.
- Wagner LK and Archer BR, Minimizing Risks from Fluoroscopic X Rays, 2nd ed., R.M. Partnership, The Woodlands, TX.
- Balter S, et al. Fluoroscopically guided interventional procedures: A review of radiation effects on patients' skin and hair. *Radiology*, 254:326-341
- NCRP Report 168