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Establishing and Using DRLs for optimisation in Radiography

Sue Edyvean

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Acknowledgment for slides from:
J.Vassileva@iaea.org, jenny.smith@phe.gov.uk

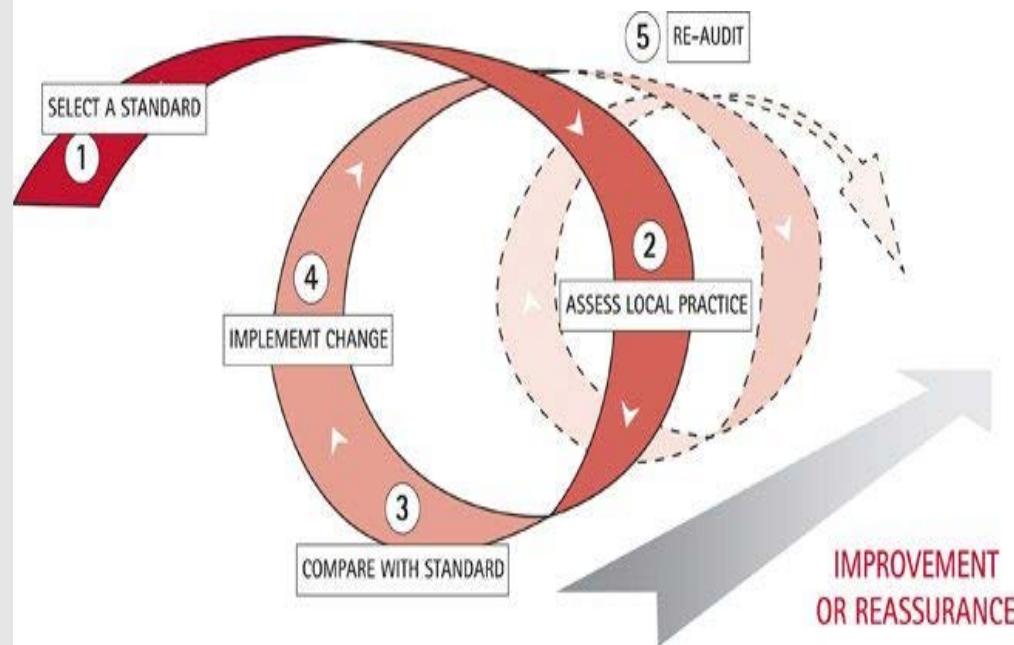
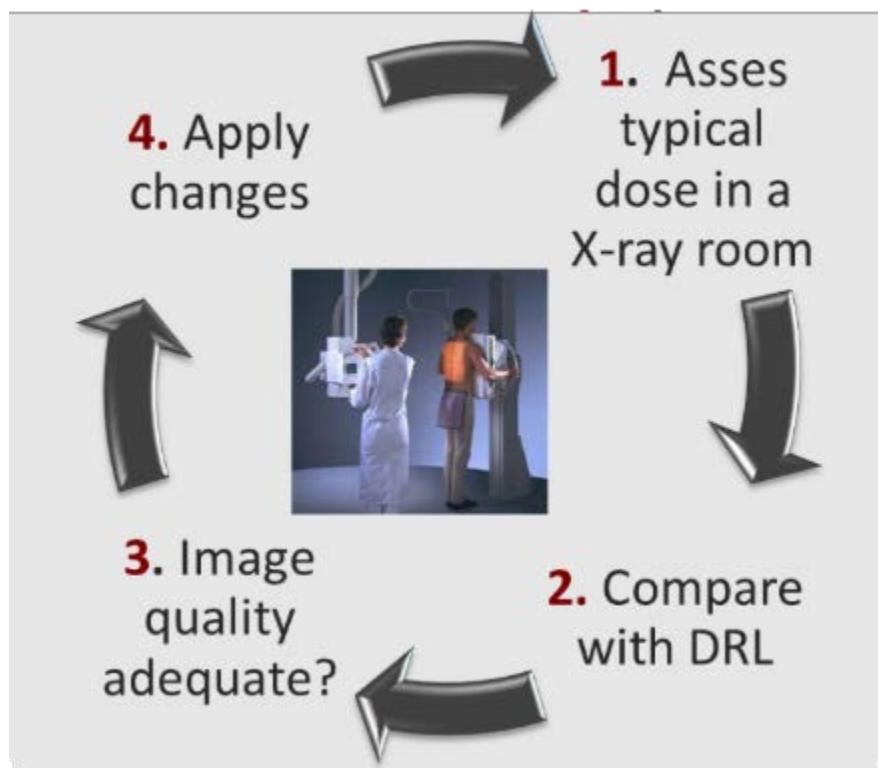
Senior Scientific Group Leader
Medical radiation Dosimetry, CRCE
Public Health England
Didcot, Oxon. OX11 0RQ, UK



Outline

- Radiology dose audit
 - Dose quantities
 - Establishing DRLs
- Optimisation
 - Radiology Equipment
 - Radiographic Practice

Patient dose audit and DRLs



The function of DRLs is to stimulate a continuous cycle of assessment and optimisation of practice

Designing the dose audit to obtain the DRL

1. Select examination
2. Select patient data
3. Ensure weight is within required range
4. Record the dose index value
5. Review and clean the data
6. Calculate the median value of dose index

Designing the dose audit to obtain the DRL

1. Select examination

- a. High dose
- b. High frequency

Radiography examinations, their relative frequencies and contribution to collective effective dose

Examination	Percentage of total frequency of all radiography examinations (%)	Percentage contribution to collective dose (%)
Chest and thorax	12-29	0.7-5.2
Abdomen, pelvis and hip	7.4-14.3	2.9-14
Spine (thoracic and lumbar)	3.8-12.7	30
Intravenous urography	0.3-2.0	1.2-8.7

- As chest radiography is a very common examination and involves exposure of several radiosensitive organs, it should be included in surveys of radiography.
- The largest contributions from radiography to collective effective dose are examinations of the abdomen, pelvis, and spine, so these should also be included in any radiographic survey.
- If performed with a sufficient frequency, it is recommended that skull X rays should be included, as they involve exposure of the lens of the eye.
- High frequency/low dose exams such as, x-rays of extremities should not necessarily be a high priority:

NHS England 2016

Planar Radiography (from RIS records)

No.	SNOMED-CT code name	Total exam count in 2016	% Total Plain Radiography
1	Plain chest X-ray	7,767,185	37.9%
2	Radiologic examination of knee	1,599,860	7.8%
3	Radiography of foot	1,014,505	4.9%
4	Pelvis X-ray	1,003,500	4.9%
5	Radiography of ankle	843,070	4.1%
6	Radiography of shoulder	778,775	3.8%
7	Diagnostic radiography of abdomen	723,920	3.5%
8	Radiography of hand	673,860	3.3%
9	Radiography of wrist	633,005	3.1%
10	Radiography of hip	632,930	3.1%
11	Diagnostic radiography of lumbar spine	432,400	2.1%
12	Radiography of cervical spine	279,615	1.4%

Selecting Projections/Exams

- First on the high dose and high frequency X-ray (red)
- Next exams of slightly lower dose or frequency (gold)
- Then low dose but very high frequency exams, and low frequency but high dose exams (yellow)
- Providing typical doses for extremity exams (White).
Is setting an NDRL appropriate?

If X-ray frequencies are given only by body region then projections and complete exams will need to be chosen.

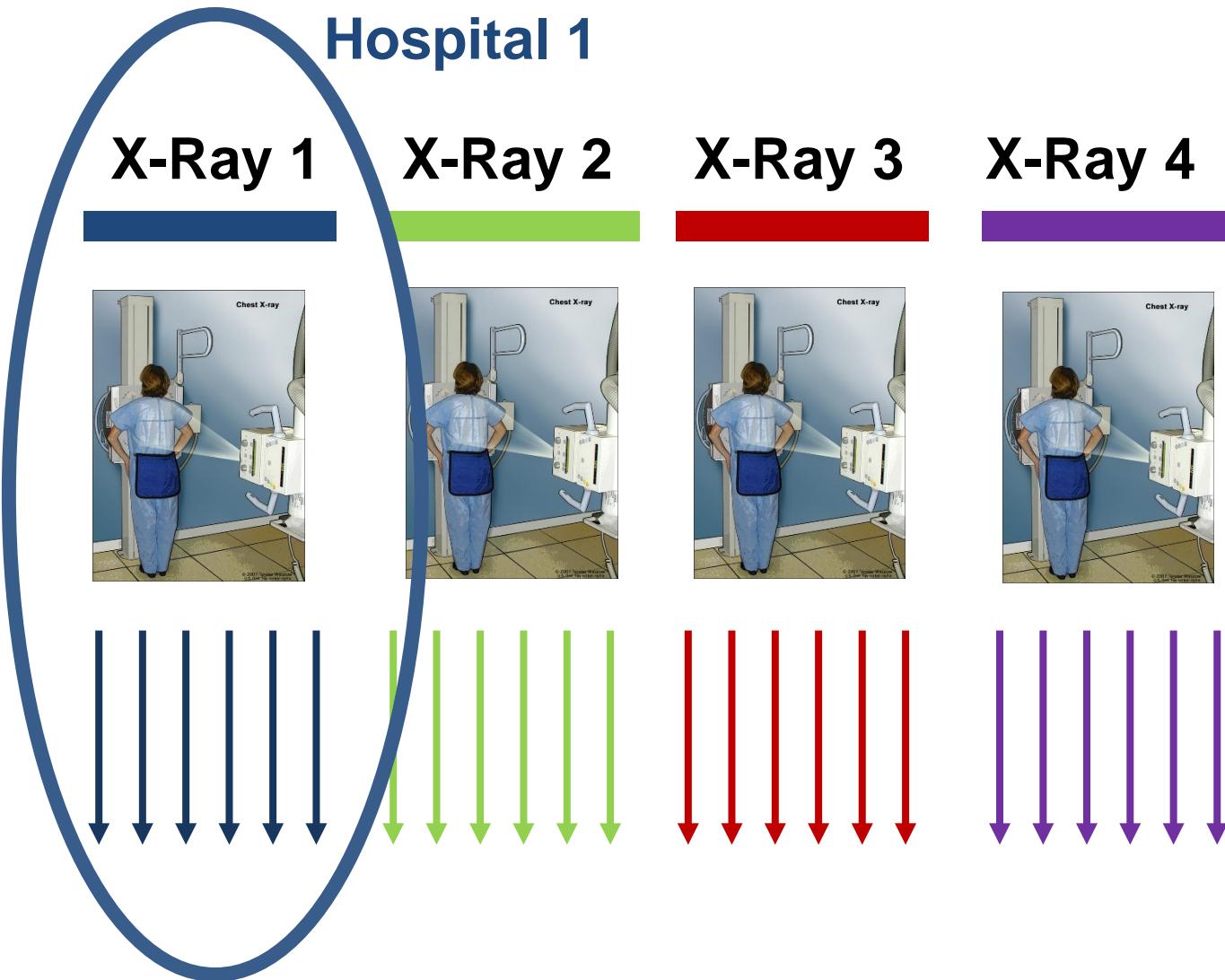
Choosing an Exam list

- Exam frequency may be determined by several routes:
e.g.
 - patient records
 - RIS records
 - National (or subregion) dataset/registry
 - Initial survey

Reasons for not setting KAP NDRL for Extremities

- Usually very low dose exposures, so a low priority
- Exposure extends beyond the edges of body part so collimation of image is a matter of choice.
- KAP is determined from beam size, and hence will vary with collimation, and overestimate the patient's (low) exposure.
- Both hands or both feet can be x-rayed in one image but records can be unclear if both extremities, or only one, was imaged. Where mass data is being retrospectively accessed it is not practical to check this visually.
- Knee KAP DRLs can be criticised for the same reasons.

Dose Audit on first x-ray room



Designing the dose audit to obtain the DRL

2. Select patient data – how many (as minimum)?
 - 20 – 30 patients
3. Ensure weight is within required range
 - ICRP 135 ? : ‘standard size’
 - Previous advice and a useful starting point
 - Patients between 50 – 90 kg (70kg +/- 20 kg)
 - Giving a mean value f 70 kg +/- 10 kg (or 5 kg)

Dose Audits – Numbers of data and patient size

- (227) If data collection is via paper forms, the number of patients will be limited, but should be at least 20–30. With restricted numbers, information on patient sizes should be recorded, if possible, or at least the range of sizes should be restricted, with very large and very small patients being excluded. This is not a concern when an automated data collection system is used.

ICRP 135

Designing the dose audit to obtain the DRL

4. Record the dose index value
5. Review the data
6. Calculate the median value of dose index

Radiology DRL Quantities

DRL quantity	Air kerma-area product	Entrance surface air kerma
Symbols	P_{KA}	$K_{a,e}$
Recommended Units for DRL [^]	mGy.cm^2	mGy
Other symbols in common use	KAP	ESAK
Closely similar quantity	Dose-Area product (DAP)*	Entrance-surface dose (ESD)*

[^] Recommended by IAEA and ICRP. Note UK uses Gy.cm^2 for DAP NDRLs

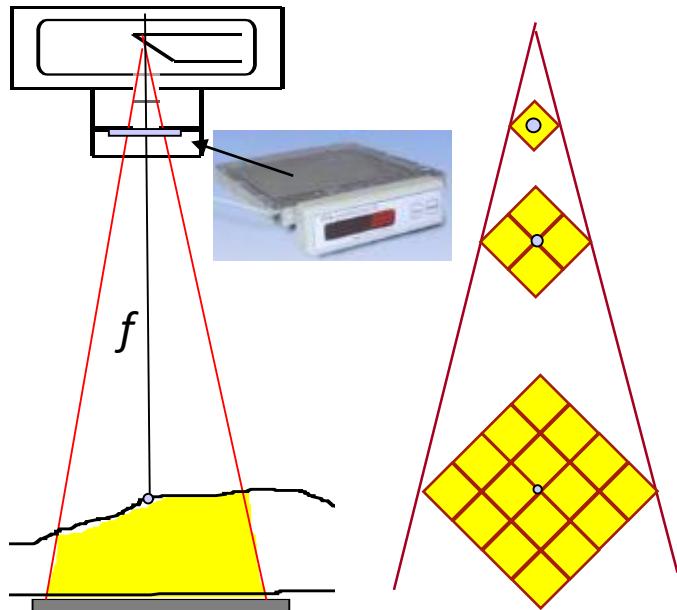
* Because "air kerma" and "dose in air" are numerically equal in diagnostic radiology energy range

1. Measurable dose quantities

Radiography and fluoroscopy

Air kerma-area product, KAP

Also known as Dose Area Product, DAP



$$\text{Kerma (dose)} \sim (1/f)^2$$

$$\text{Field area} \sim f^2$$

KAP is independent on distance from focus

Unit: $\text{Gy} \cdot \text{m}^2$

Practical units:

$$1 \mu\text{Gy} \cdot \text{m}^2 = 1 \text{ cGy} \cdot \text{cm}^2$$



KAP dose units

Radiology system may display KAP doses in many units:

Gy.cm²

dGy.m²

μ Gy.cm²

cGy.cm²

cGy.m²

mGy.cm²

mGy.m²

dGy.cm² (can be displayed as dGy.cm² or DGYCM2)

μ Gy.m² (can be displayed as μ GY.m² or as μ GYM2)

- Note that 1 cGy.cm² and 1 μ Gy.m² are the same, 1/100 of 1 Gy.cm².

Machine displays may not use lower case letters, and so display mGy as MGY.

1. Measurable dose quantities

Radiography and fluoroscopy

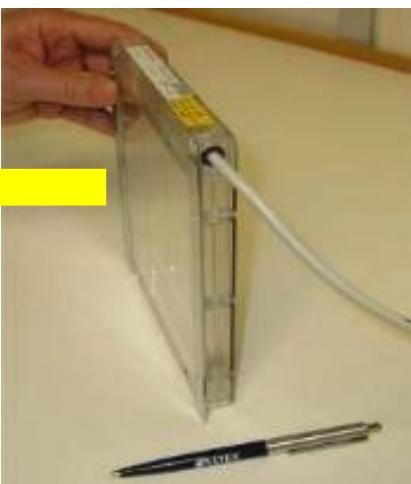
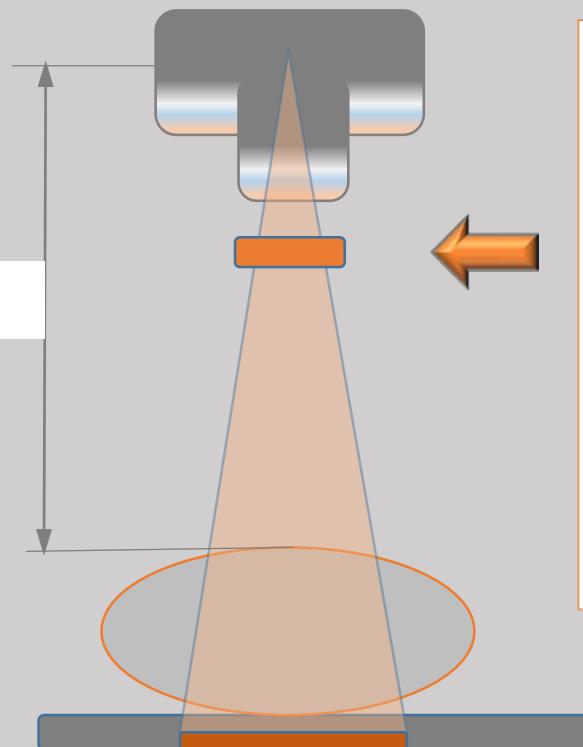


Table 1: KAP Dose Units conversion Table

If your value is currently in the original units and has a value of 1, the table gives its value in the new units (those you want to convert it into).

New Units	$\mu\text{Gy m}^2$							
	Gy cm^2	dGy cm^2	cGy cm^2	mGy cm^2	Gy m^2	dGy m^2	cGy m^2	mGy m^2
Original units								
$\mu\text{Gy m}^2$	Gy cm^2	1	10	100	1,000	1.E-04	0.001	0.01
	dGy cm^2	0.1	1	10	100	1.E-05	1.E-04	0.001
	cGy cm^2	0.01	0.1	1	10	1.E-06	1.E-05	0.001
	mGy cm^2	0.001	0.01	0.1	1	1.E-07	1.E-06	0.0001
	Gy m^2	10,000	1,000	100	10	1	10	100
	dGy m^2	1,000	10,000	1.E+05	1.E+06	0.1	1	10
	cGy m^2	100	1,000	10,000	1.E+05	0.01	0.1	1
	mGy m^2	10	100	1,000	10,000	0.001	0.01	0.1

Air kerma-area product



Air kerma-area product (P_{KA})

$$P_{KA} = K \cdot A$$

K – air kerma

A – area

Unit: Gy·m², mGy·cm²,
cGy·cm², μ Gy·m², Gy·cm² ...



Air kerma-area product

Air kerma-area product (P_{KA})

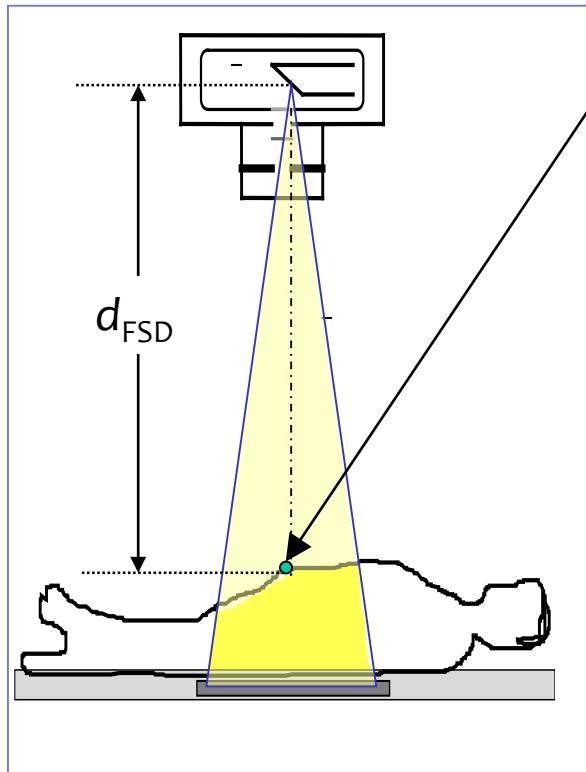
P_{KA} values recorded by meters, calculated by the equipment, or given by the manufacturers and reported in the DICOM header should be reasonably accurate. Arrangement should be in place to check the calibration of P_{KA} meters and the accuracy of P_{KA} values calculated and displayed by the X ray equipment and recorded in the DICOM header.

Note: Standard IEC 60580 specifies acceptable limits of uncertainty in the response of P_{KA} meters when individual exposure vary to the maximum likely extent. According to this, the estimated expanded uncertainty of a measurement with P_{KA} meters is 25% at the 95% confidence interval ($k = 2$).

References: IAEA TRS 457, IAEA, Vienna, 2007 and Lin, PJ, et. al. Accuracy and calibration of integrated radiation output indicators in diagnostic radiology: A report of the AAPM Imaging Physics Committee Task Group 190, Med Phys, 42: 6815-6829 (2015)

1. Measurable dose quantities

Radiography

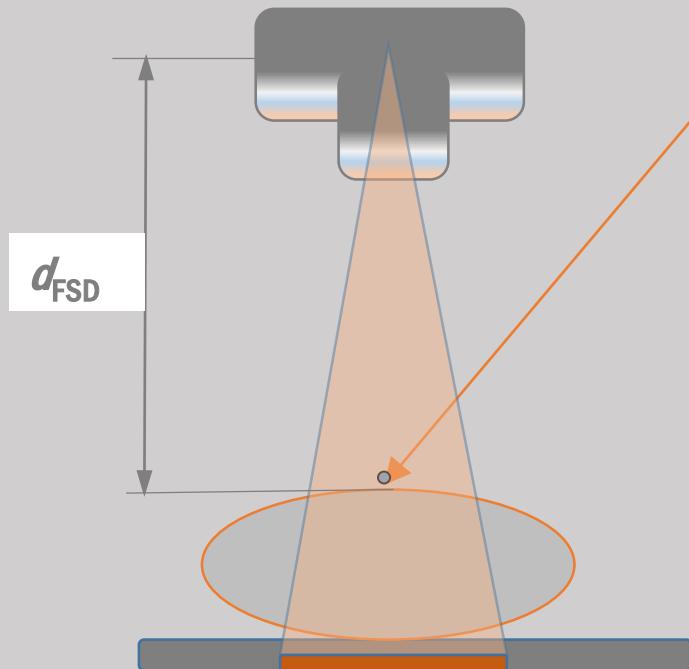


Entrance surface air kerma
 K_e [Gy]

Also known as
Entrance surface dose, ESD

- 1) Measured with phantom
- 2) With patients:
 - Directly measured with TLD
 - Calculated from the tube output

Entrance surface air kerma



Entrance air kerma, $K_{a,e}$

$$K_{a,e} = K_{a,i} \cdot B$$

B – Backscatter factor (1.2 – 1.4)

Unit: Gy

Direct measurement

Calculation from the tube output

IAEA Radiology Survey Workbook ESAK guidance

Instructions for calculation of entrance surface air kerma from the x-ray tube output

Definitions

Incident air kerma is the air kerma from the incident beam on the central x-ray beam axis at the focal-spot-to-surface distance at the skin entrance plane. Only primary radiation incident on the patient or phantom and not backscattered radiation is included.

Entrance-surface air kerma (ESAK) is the air kerma on the central X ray beam axis at the point where X ray beam enters the patient or phantom. The contribution of the backscattered radiation is included.

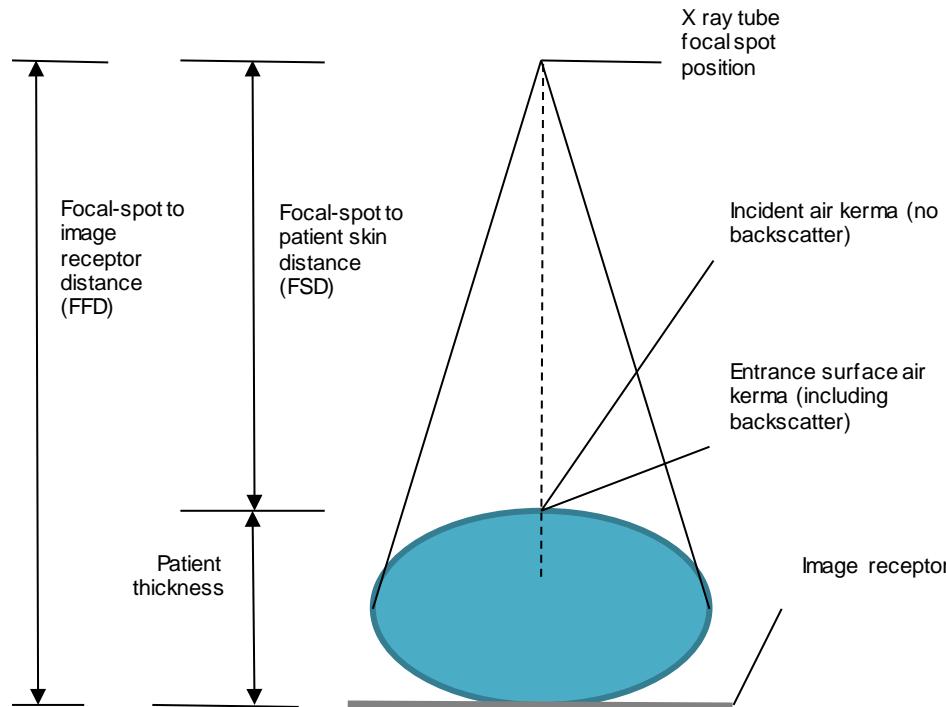


Figure 1. Schematic diagram showing some dosimetric and geometric quantities

ESAK can be established indirectly from incident air kerma using appropriate backscatter factor. Incident air kerma is estimated indirectly for the x-ray tube output at the selected distance and exposure parameters using the inverse square law.

The X ray tube output is air kerma at specified distance from the x-ray tube focal spot, divided by the I_t (tube-current exposure-time product, in units mAs). It is calculated for the particular tube voltage (U) value.

ESAK units

Radiology machine may display ESAK doses in the following units:

- Gy = 1000 mGy
- dGy = 100 mGy
- cGy = 10 mGy
- mGy
- μ Gy = 0.001 mGy

Machine displays may not use lower case letters, and so display mGy as MGY.

Designing the dose audit to obtain the DRL

1. Select examination
2. Select patient data
3. Ensure weight is within required range
4. Record the dose index value
5. Review and clean the data
6. Calculate the median value of dose index

ID	AP Projecti on DAP dose *	Patient <u>weight</u> (kg)
1	22	114.3
2	5	50.8
3	9	139.7
4	3	88.9
5	6	44.45
6	7	82.55
7	6	49
8	5	49
9	9	70
10	8	50.8
11	11	75
12	9	68
13	6	57
14	7	65
15	8	70
16	7	70
17	6	82.55
18	11	48
19	11	60
20	9	85.73
21	12	76
22	5	50.8
23	27	114.3
24	7	76.2
mean	9	72.42
median	7.5	70

Designing the dose audit to obtain the DRL

1. Select examination
2. Select patient data
3. Ensure weight is within range
4. Record the dose index value
5. Review and clean the data
6. Calculate the median value
- “Typical value”

Large and small patients removed

Record ID	AP Projection DAP dose *	Patient weight (kg)
2	5	50.8
4	3	88.9
5	6	44.45
6	7	82.55
9	9	70
10	8	50.8
11	11	75
12	9	68
13	6	57
14	7	65
15	8	70
16	7	70
17	6	82.55
18	11	48
19	11	60
20	9	85.73
21	12	76
22	5	50.8
23	27	114.3
24	7	76.2
mean	7.6	68.0
median	7.0	70.0

ID	AP Projection DAP dose *	Patient weight (kg)
1	22	114.3
2	5	50.8
3	9	139.7
4	3	88.9
5	6	44.45
6	7	82.55
7	6	49
8	5	49
9	9	70
10	8	50.8
11	11	75
12	9	68
13	6	57
14	7	65
15	8	70
16	7	70
17	6	82.55
18	11	48
19	11	60
20	9	85.73
21	12	76
22	5	50.8
23	27	114.3
24	7	76.2
mean	9	72.42
median	7.5	70



Mean versus Median – Simple tutorial



Mean versus Median

Mean	Median
Average of values	Same number of data points above and below (50 th percentile)
More affected by outliers	Less affected by outliers
Less robust for skewed distributions	More robust for skewed distributions

Nine numbers: 7 9 11 6 13 6 6 3 11			
Put in order	3 6 6 6 7 9 11 11 13	Mode	6
Put in order	3 6 6 6 7 9 11 11 13	Median	7
Add all	$7+9+11+6+13+6+6+3+11 = 72$ There are 9 numbers: $72 \div 9 = 8$	Mean (average)	8



Mean versus Median

Mean	Median
Average of values	Same number of data points above and below (50 th percentile)
More affected by outliers	Less affected by outliers
Less robust for skewed distributions	More robust for skewed distributions

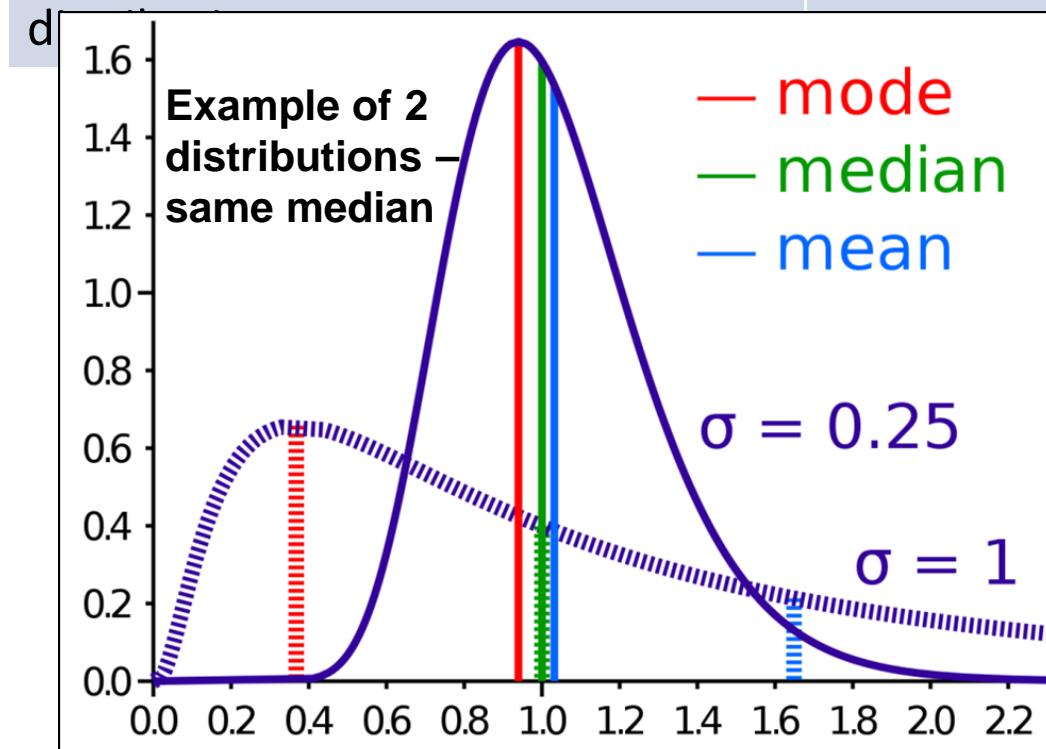
If highest value is 130 not 13:

Nine numbers: 7 9 11 6 <u>130</u> 6 6 3 11			
Put in order	3 6 6 6 7 9 11 11 <u>130</u>	Mode	6
Put in order	3 6 6 6 7 9 11 11 <u>130</u>	Median	7
Add all	$7+9+11+6+\underline{130}+6+6+3+11 = 189$ There are 9 numbers: $189 \div 9 = 8$	Mean (average)	21



Mean versus Median

Mean	Median
Average of values	Same number of data points above and below (50 th percentile)
More affected by outliers	Less affected by outliers
Less robust for skewed distributions	More robust for skewed distributions



Nine numbers: 7 9 11 6 13 6 6 3 11	Mode	6
	Median	7
= 72 ÷ 9 = 8	Mean (average)	8

Analysis: in a particular X-ray room

- For the most frequently performed exams
 - For a particular age/weight group
 - Register dose values for a sample of 20-50 patients
 - Calculate median/ average dose index
 - **Typical dose value = median of the sample**



Chest X-ray
Adult patient (70 kg)
Range (Min-Max):
(0.01 – 0.10) Gy.cm²
Median = 0.1 Gy.cm²



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Analysis in a Hospital

Hospital 1

X-Ray 1



X-Ray 2



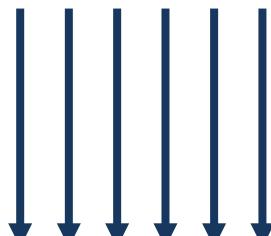
X-Ray 3



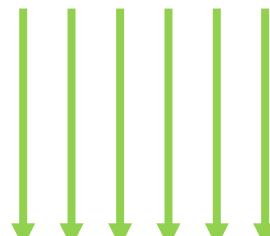
X-Ray 4



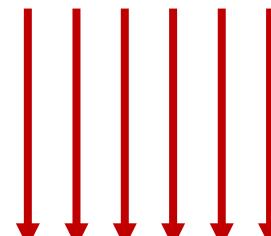
.... etc



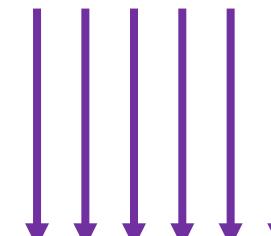
Median



Median



Median



Median

Analysis: in a hospital

Assess typical dose values for all equipment performing the same exams

Chest X-ray, Adult patient (average weight 70 kg)

Room 1:

Median = 0.1 Gy.cm²



Room 2:

Median = 0.05 Gy.cm²



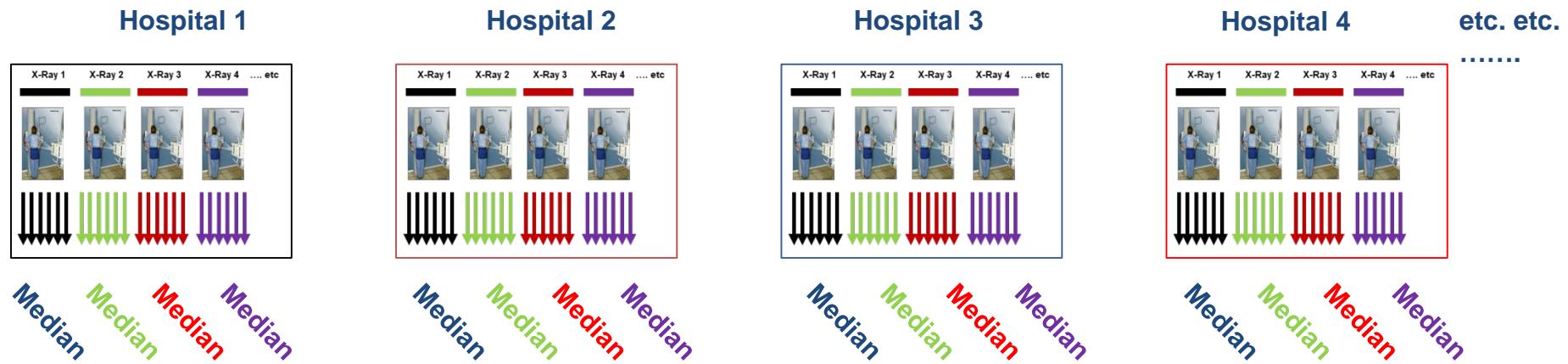
Room 2:

Median = 0.2 Gy.cm²

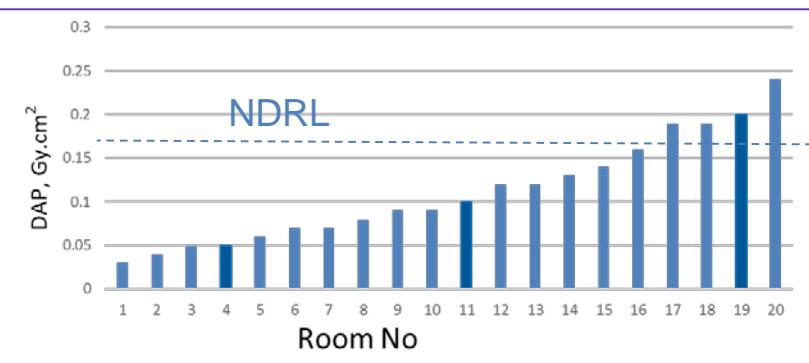


Analysis in a Country / Region - National DRLs

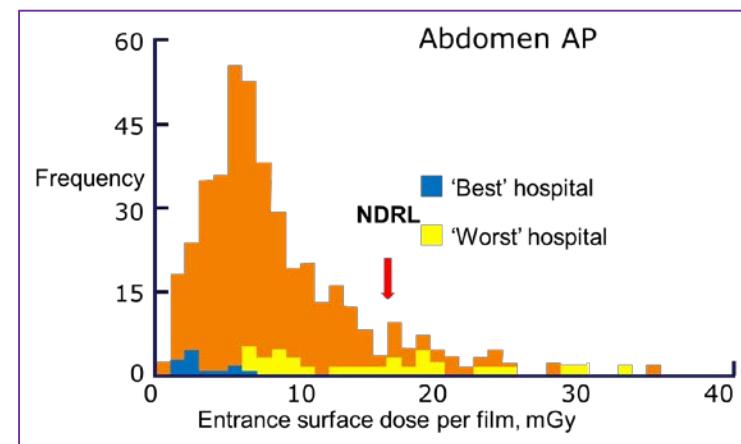
- Dose index measured for each patient; median for each system obtained:



- Show data in two ways (DRL shown in each case – 3rd quartile):



a) Room versus increasing dose index (DAP in this example)

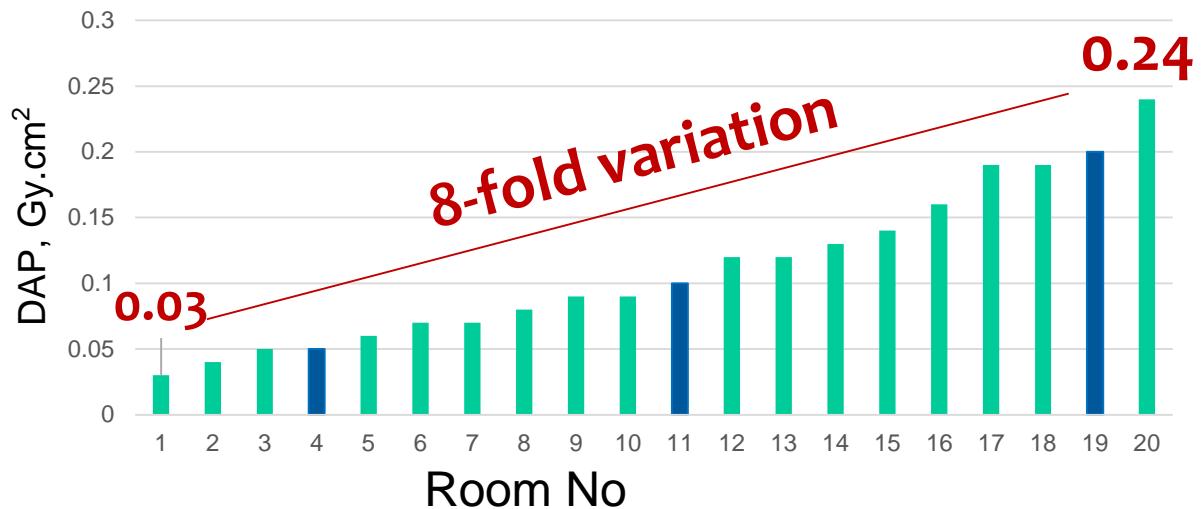


b) No. of rooms for a given dose index bin (ESD in this example)

Analysis: in a country/ region

Collect data from a representative sample of hospitals performing the same exam

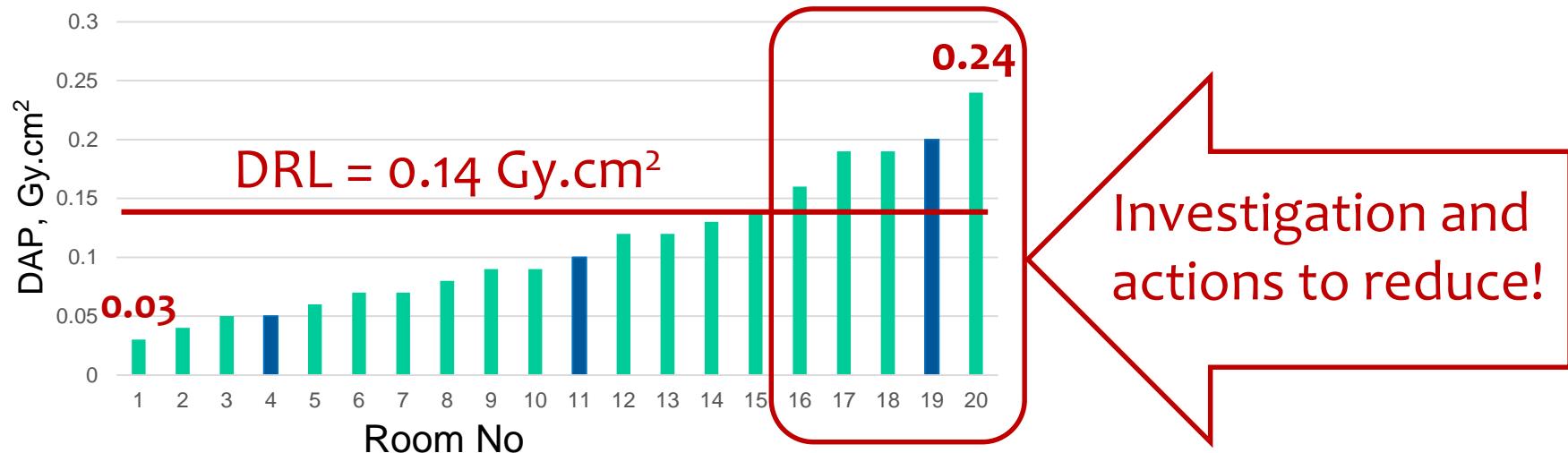
Chest X-ray, Adult patient (70 kg)
Typical DAP values in 20 X-ray rooms



Analysis: in a country/ region

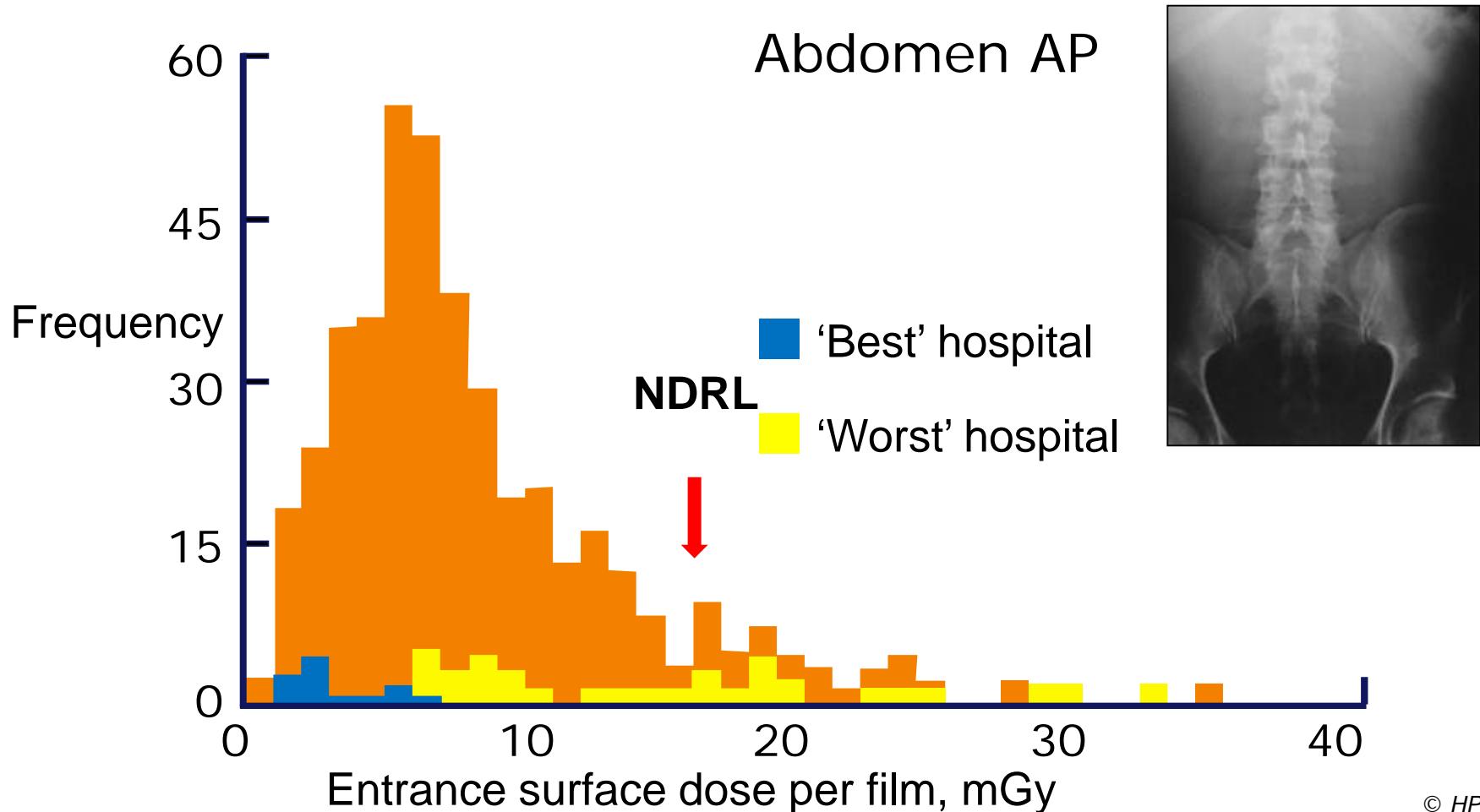
**Establish Diagnostic Reference Level (DRL)
At 3rd Quartile of dose distribution**

Chest X-ray, Adult patient (70 kg)

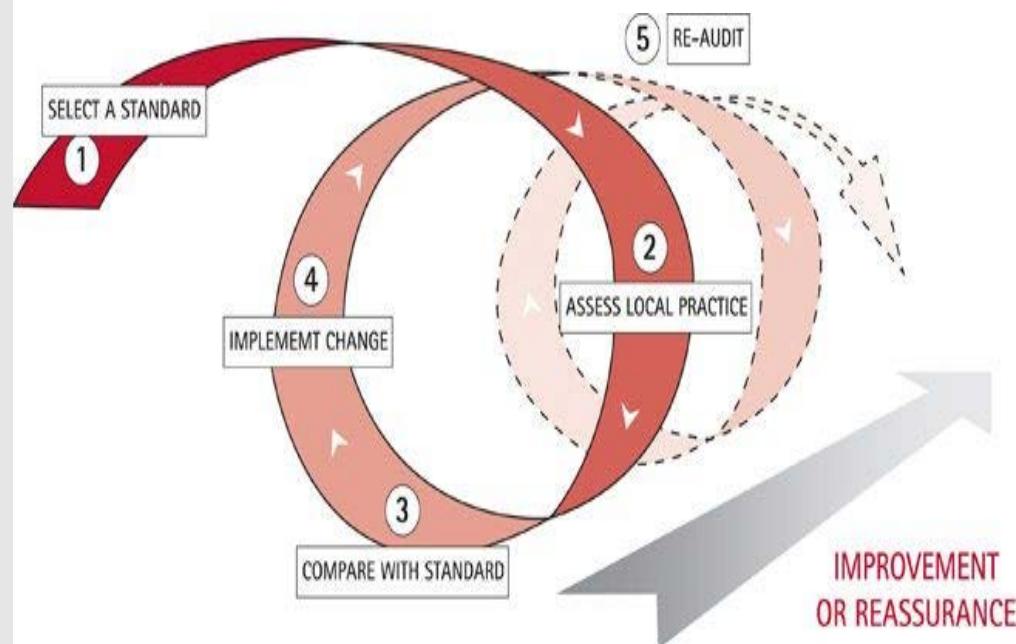
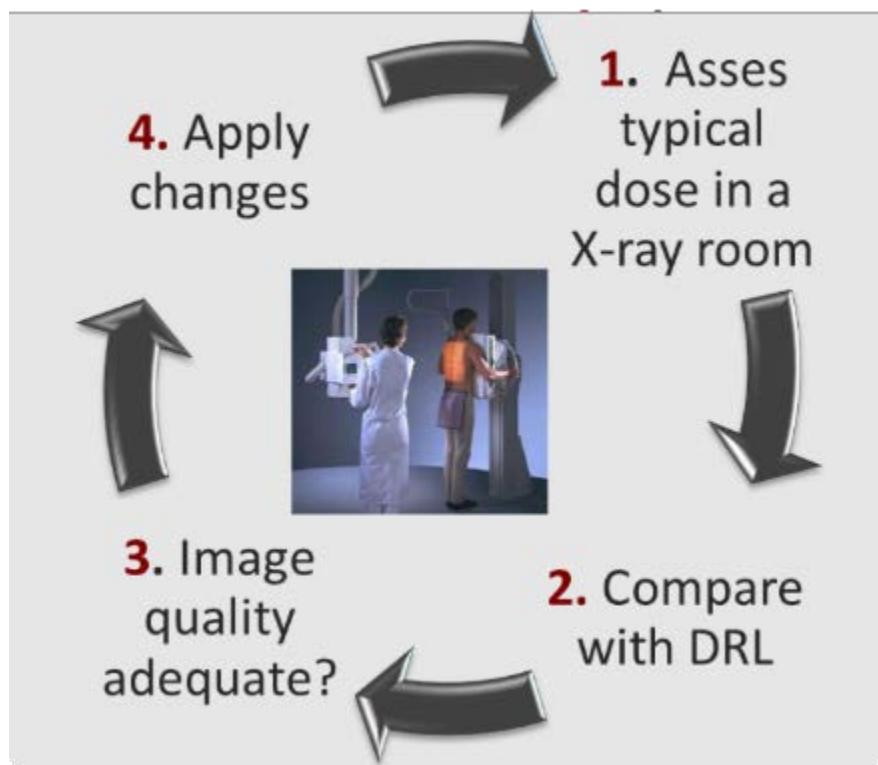


National Reference Dose – National DRL

Early NRPB Patient Dose Survey (England)



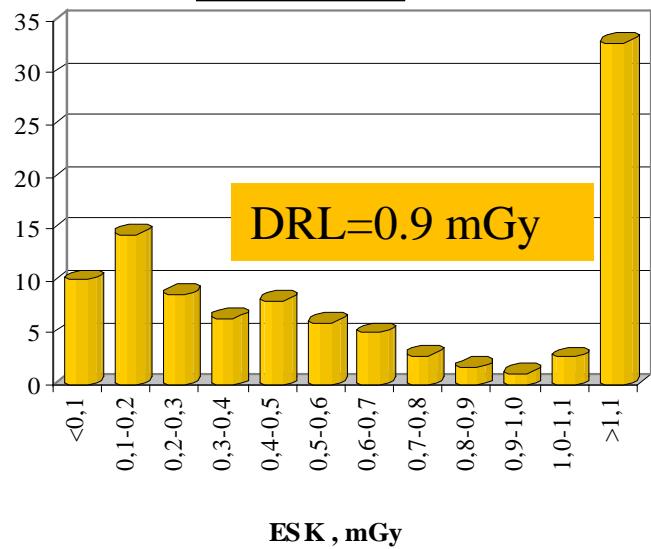
Patient dose audit and DRLs



Patient doses - radiography

National surveys in Bulgaria, Chest PA

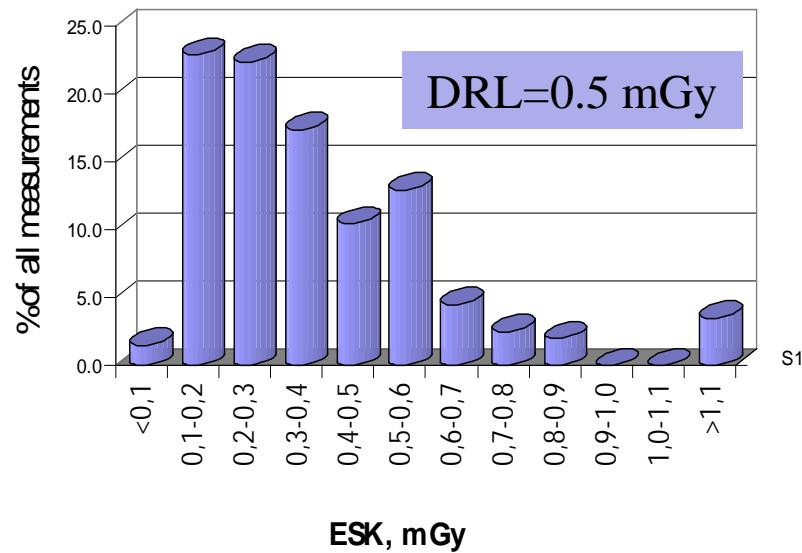
2002-2004



Mean doses: 0.06 – 2.3 mGy (**35 fold**)

Individual doses: 0.03 – 9.3 mGy

2008-2009



Mean doses: 0.1 – 1.3 mGy (**13 fold**)

Individual doses: 0.07 -3.2 mGy



Optimising

- Know the Technology
- Look to implement good radiographic practice



Equipment: Detectors Type

- **Film**
 - Film specification: make, type, speed class, size...
- **Computer Radiography (CR) Storage Phosphor techniques**
 - Generator settings and dose information not stored on CR panel
- **Flat Panel Detectors (DR)**
 - May use direct or indirect x-ray conversion to electrical signal
- **Charge-coupled device systems**
 - Small share of market; image of luminous screen recorded by a charged couple device camera and converted into a digital image
- **Photon-counting detector**
 - Excellent efficiency and other advantages. Beginning to be used for CT and digital radiography. Being used in mammography.

Radiographic imaging chain

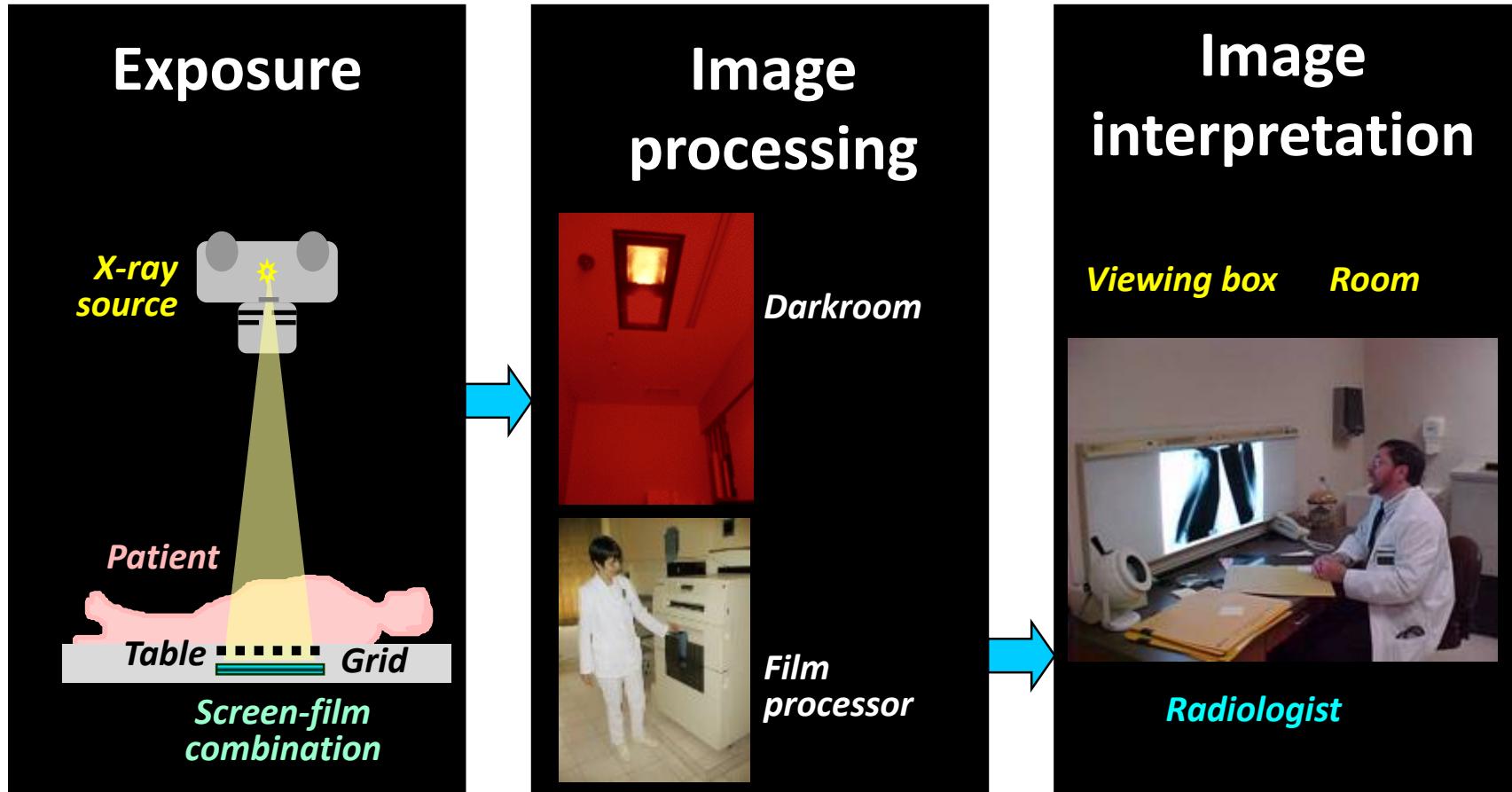


Image formation in radiography





Equipment: Digital Detectors

- Digital detector systems have high dynamic range
 - Higher doses provide higher quality images without the saturation experienced with film based system.
 - This means that there is not an automatic practical limitation to increasing patient dose as when using film.
 - QA and audit programmes required to ensure optimisation of doses to clinical tasks and to prevent “dose creep”.
- Digital detectors tend to be more sensitive than film
 - Protocols and doses should be set appropriate to the technology being used.
- Different detector technologies have different energy detection efficiency profiles
 - different energy beams and filtration may be used.



Equipment/Protocol

Other information that it may be useful to have includes to determine why doses from different sources differ:

- Machine settings (see next slide)
- Detector arrangement... Is the film/detector directly in contact with the patient or positioned in a bucky (holder) under the bed or in a support frame?
- Field size

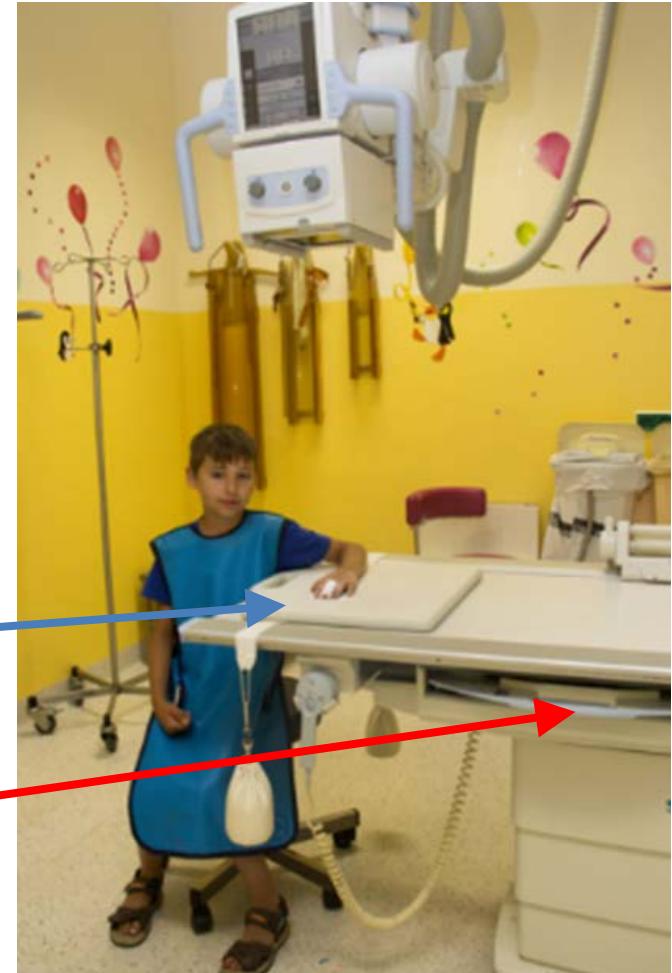


Image formation in radiography

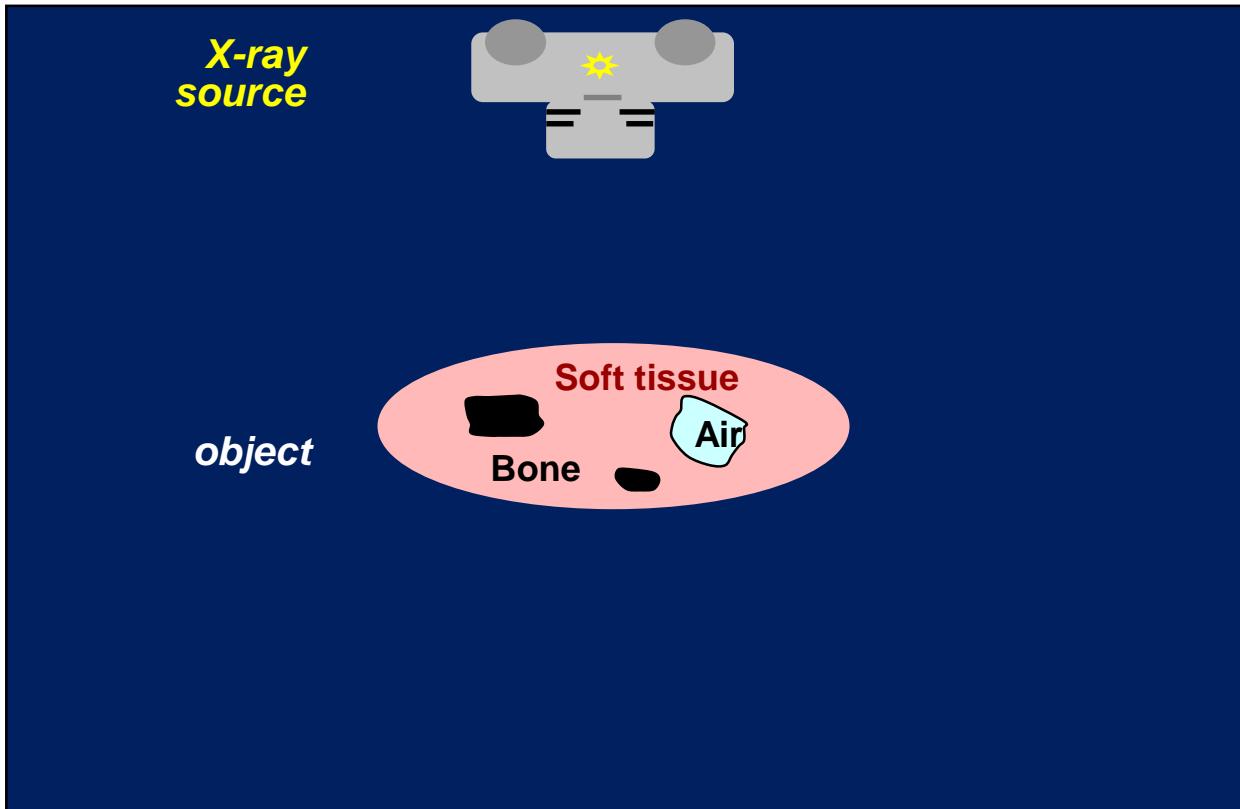


Image formation in radiography

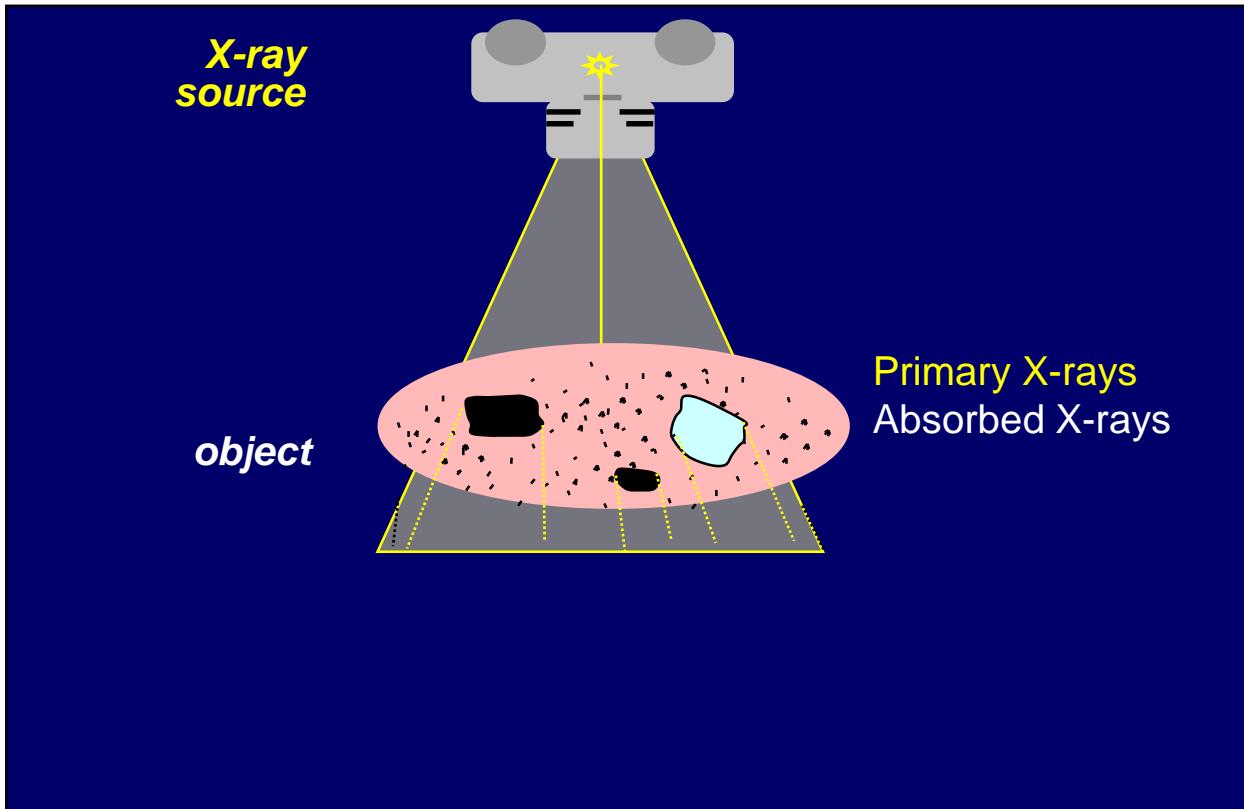


Image formation in radiography

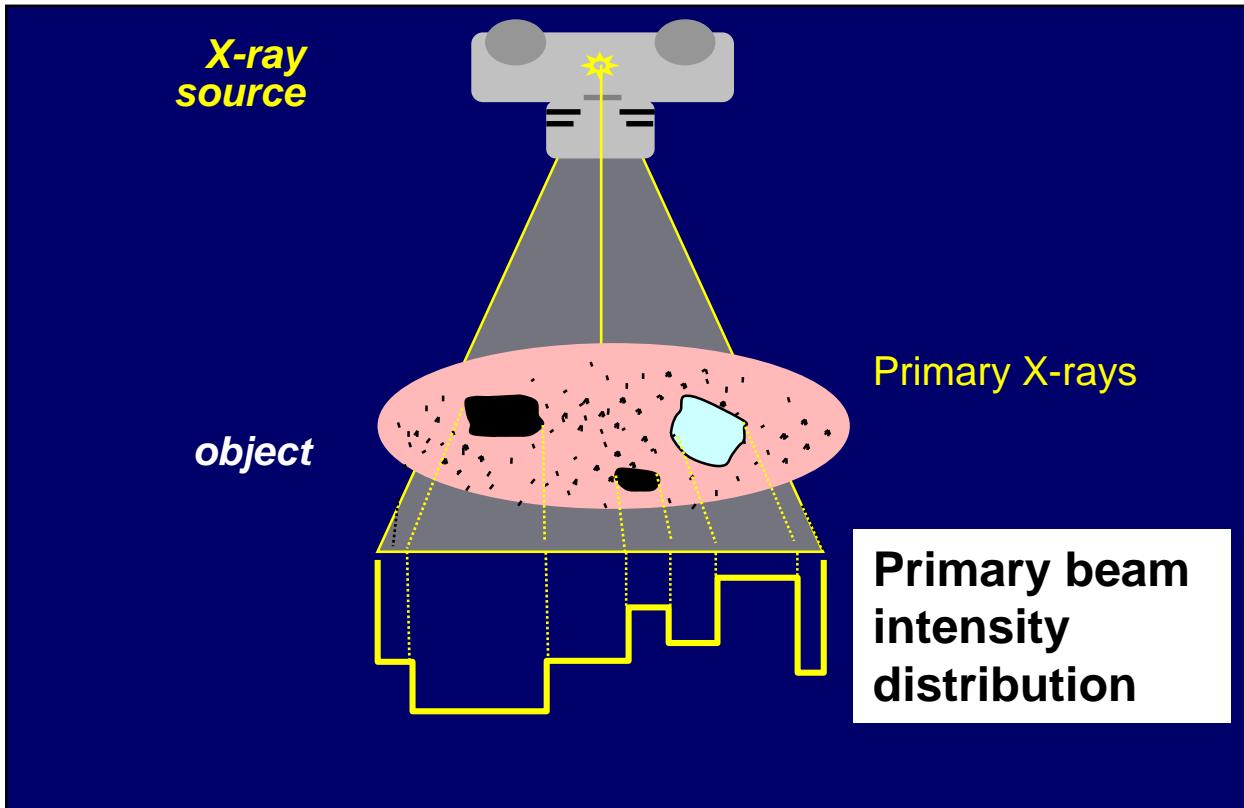


Image formation in radiography

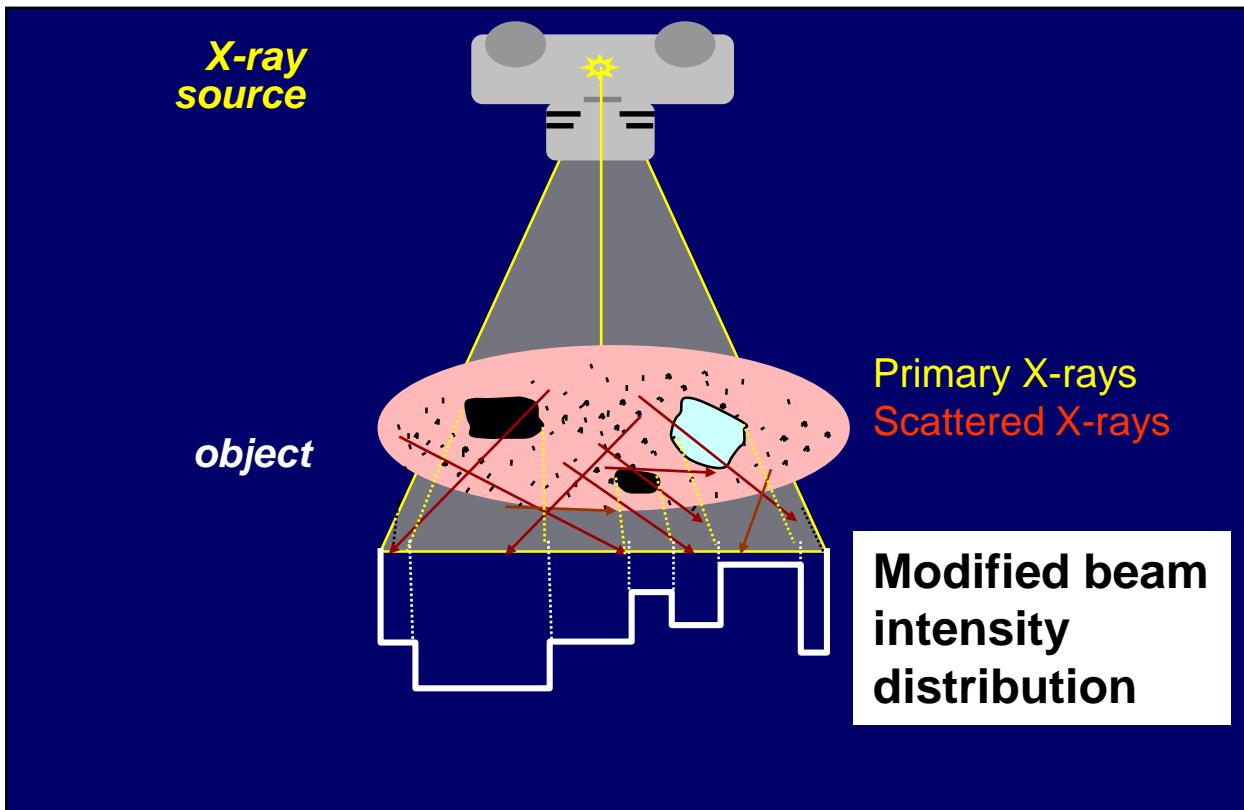
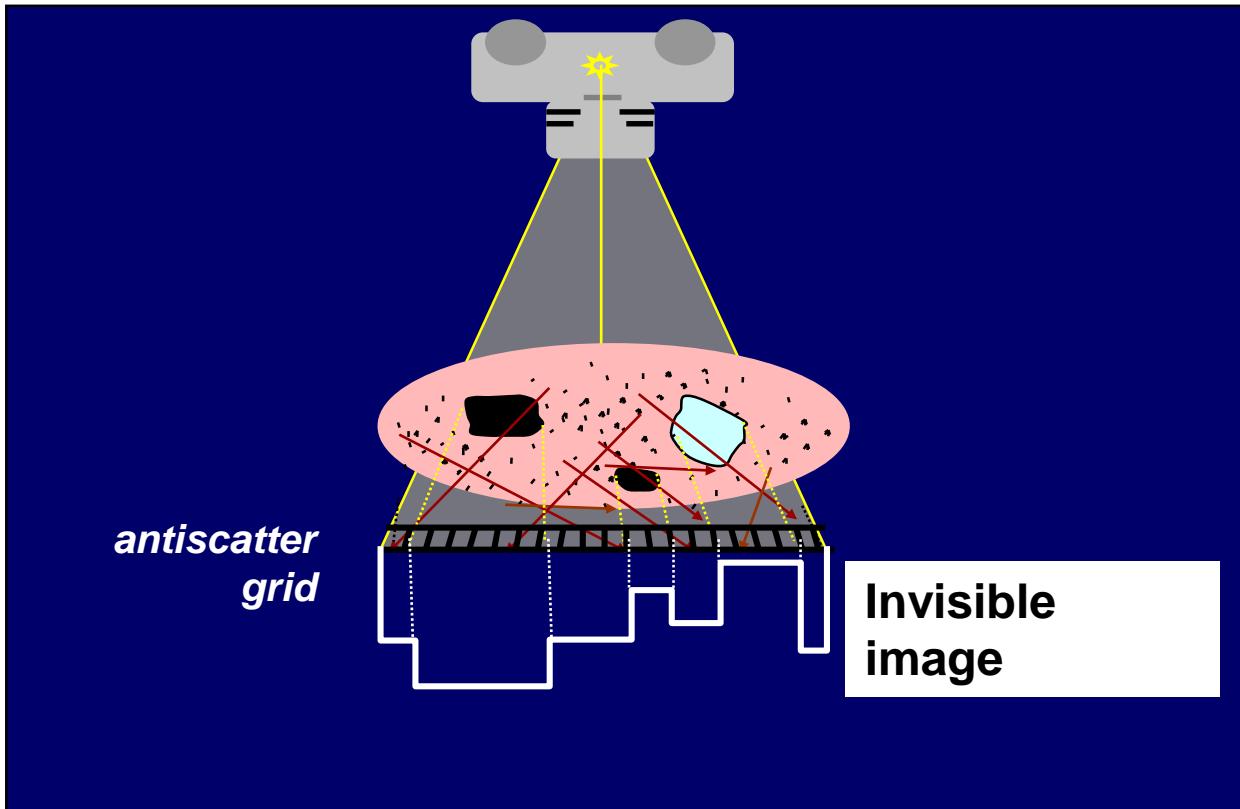
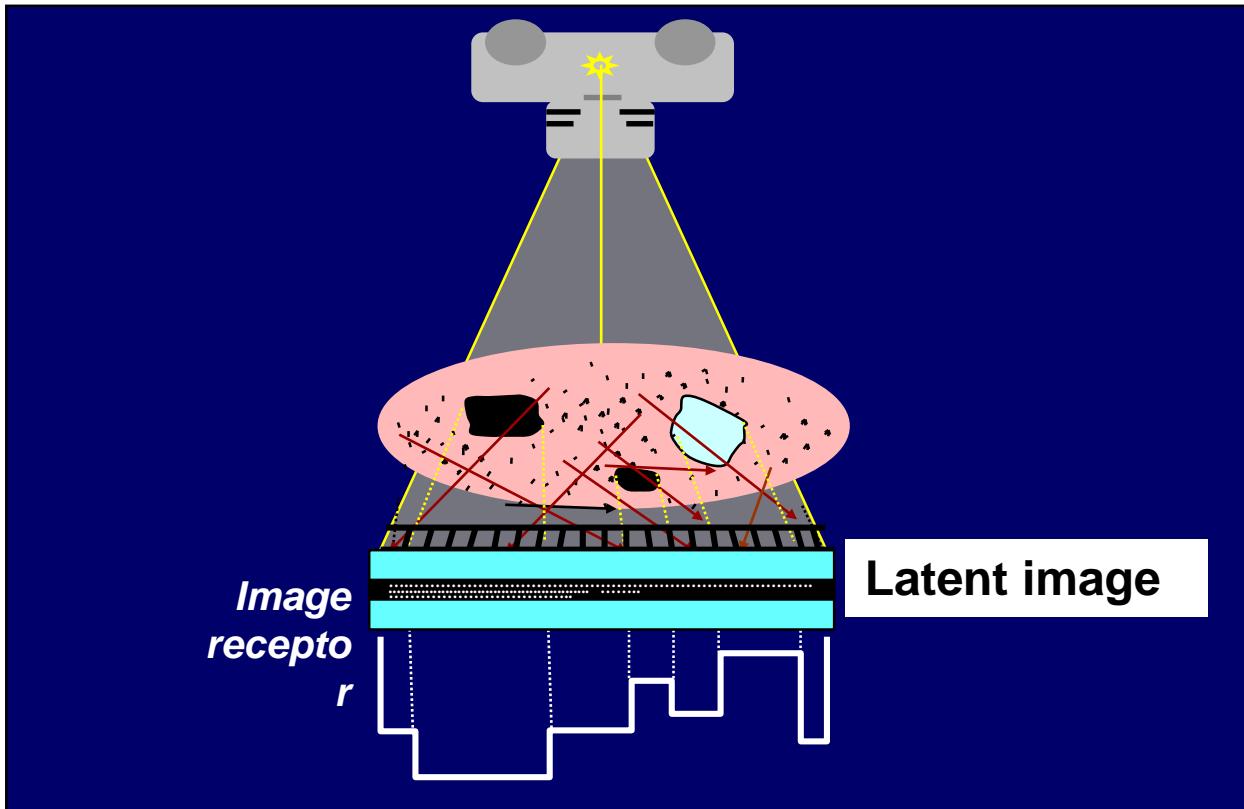


Image formation in radiography



The purpose of the collimator

Image formation in radiography



Optimization: Film-screen radiography

- Projections (AP, PA, Lat)
- Patient positioning, immobilisation / compression
- Technical parameters:
 - The tube potential (kV)
 - Tube current (mA)
 - Exposure time
 - Focal spot size
 - Filtration
 - Source to image receptor distance
 - Choice of anti-scatter device
 - Collimation
 - Image receptor type and size
- Shielding of radiosensitive organs when appropriate

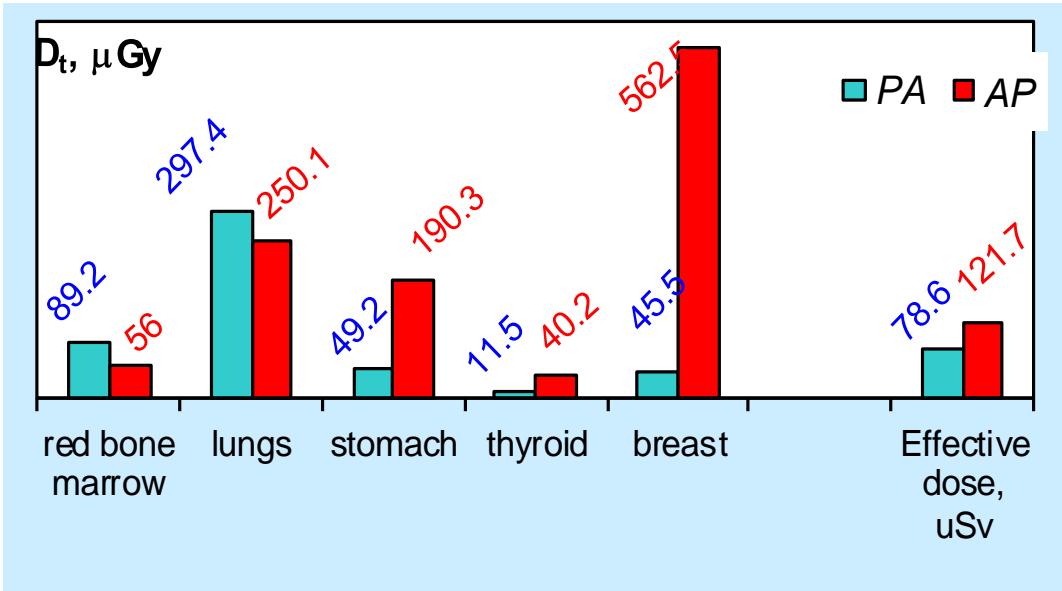


Radiography

Selection of projection

- The most appropriate projection should be adopted, from the patient protection point of view, when the diagnostic information so allows.

- ✓ For pregnant women, PA abdominal projections are preferable to minimize uterus dose
- ✓ For skull examinations eye lenses are better protected in PA projection
- ✓ For chest radiography, PA projection



Radiography

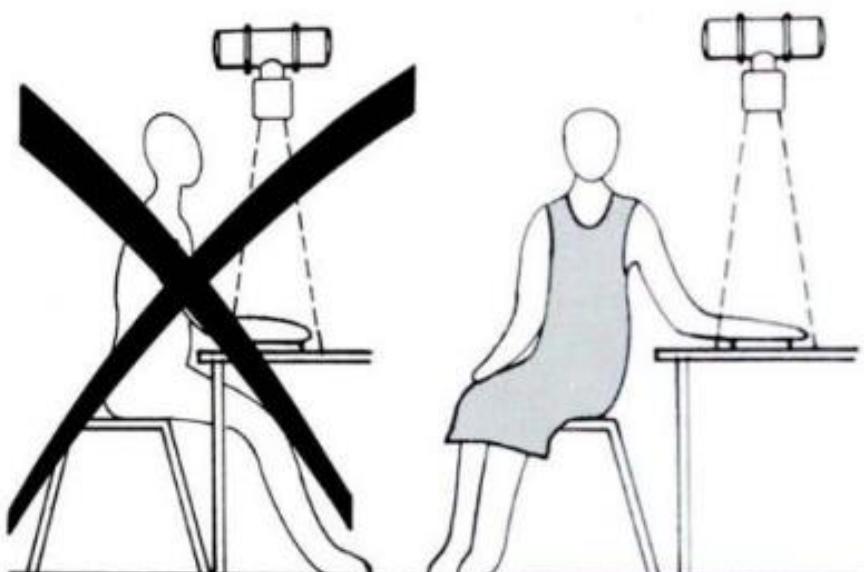
Patient positioning

- Plays a major role in determining the success of the examination
- Routine positioning may need to be altered in the light of specific clinical circumstances
- Responsibility of the person who is physically directing the examination.
- Use of suitable immobilization techniques
- **The use of fluoroscopy for centering the radiation field as a preliminary step of the radiographic image is not considered as a good radiological practice, so that it should be avoided.**



Radiography

Patient positioning



Courtesy E. Sorantin

Radiography

Patient immobilization and protective shielding



Courtesy D. Koff

Courtesy E. Sorantin

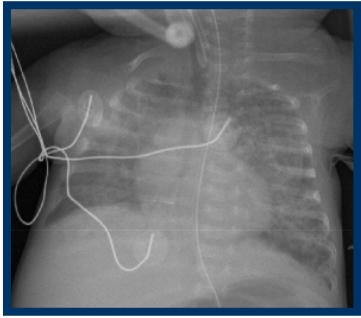
Radiography

Collimation

- The most important factor for improving image quality whilst also reducing dose
- The most common radiographic fault
- **Good collimation/coning** is essential to achieve **better contrast** and avoid exposing unnecessarily other body parts (**dose reduction**)
- Body parts outside the region of interest should not be in the X-ray field

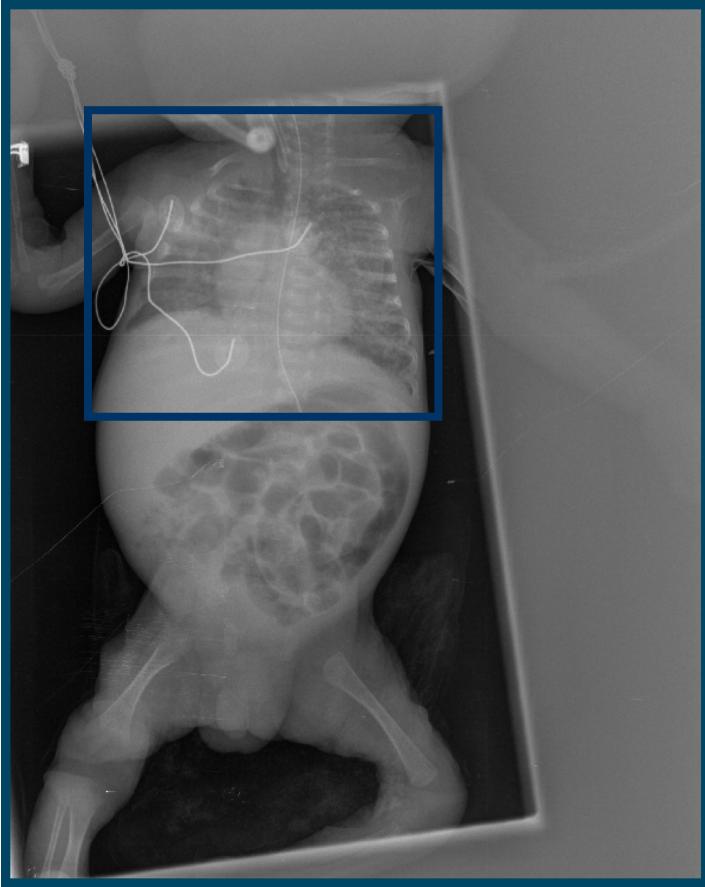


Radiography



- Poor collimation
- Large part of the body is irradiated
- Not seen on digitally cropped image

Radiography



- Poor collimation
- Large part of the body is irradiated
- Not seen on digitally cropped image



Radiography

- Example. Neonatal chest radiography (real case from my practice) (Jenia)

Table, $f=100$ cm, 65kV,
13mAs, 3,5mm Al

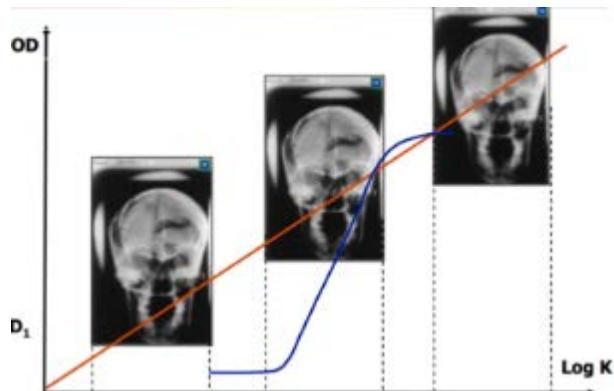


Field size	12x12 cm	20x35 cm
<i>ESD</i>	820 μGy	820 μGy
<i>ovaries</i>	26	525
<i>testicles</i>	1,6	770
<i>red bone marrow</i>	67	201
<i>lung</i>	433	491
<i>stomach</i>	348	532
<i>thyroid</i>	502	743
<i>large intestine</i>	27	537
Effective dose	209 μSv	505 μSv

Radiography

Image receptor

- For analogue image acquisition systems: the type (speed and spectral response) of film–screen combination and film processing conditions (e.g. the chemicals used and developing time and temperature).
- Digital systems can be operated at almost any level of receptor dose: Receptor dose needs to be tailored to the imaging task (decision is made by a responsible radiologist)



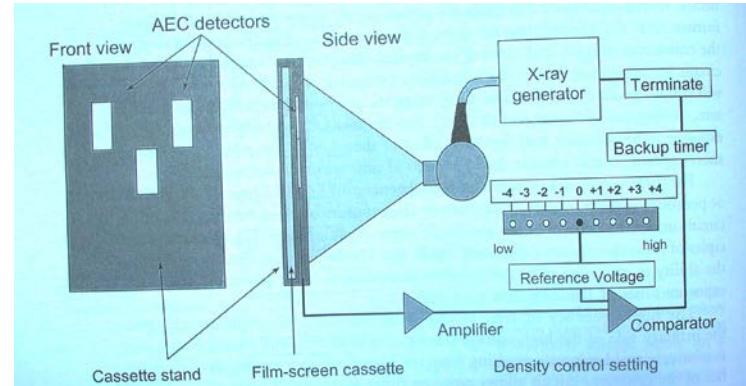
Radiography

Anti-scatter device

- Anti-scatter grid **improves the image quality**,
- but **ALWAYS increases patient doses**.
- It is advisable to evaluate whether the grid is actually necessary in equipment where its use is optional according to procedure (e.g. mammography) or patient characteristics.
- For infants and younger children the use of a grid or other anti-scatter measures is often **unnecessary**. The examples for good radiographic technique specify when grids are unnecessary.

Radiography

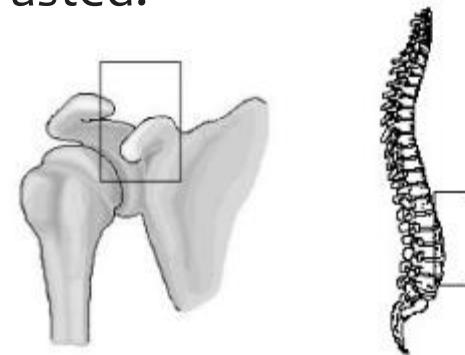
- Suitably calibrated and maintained **automatic exposure controlled systems (AEC)** should be used when available and appropriate.
- AEC systems are calibrated based on the radiation exposure at the detector required to produce the desired level of optical density (OD) for film-screen systems or a pre-determined acceptable level of signal to noise ratio (SNR), or surrogate, for digital systems.
- For digital systems, users need to understand how the selection of “exposure index” (or similar term for exposure indicator) affects patient dose.



Radiography

Automatic Exposure Control

- Inappropriate selection of automatic exposure control settings might lead to incorrectly contrasted images (too dark or too clear).
- Then the automatic exposure control device should **ALWAYS** be checked, in particular when the sensitivity of the screen-film combination has been changed.
- The correct operation of the AEC device requires, for each projection, the selection of the chamber or detector closer to the area of interest, so that this area be best contrasted.
- Generally not appropriate for small children



Radiography

Exposure parameters (example from the First survey in Bulgaria)

Example: the same type of x-ray system in 8 facilities

Chest radiography, PA, standard adult patient

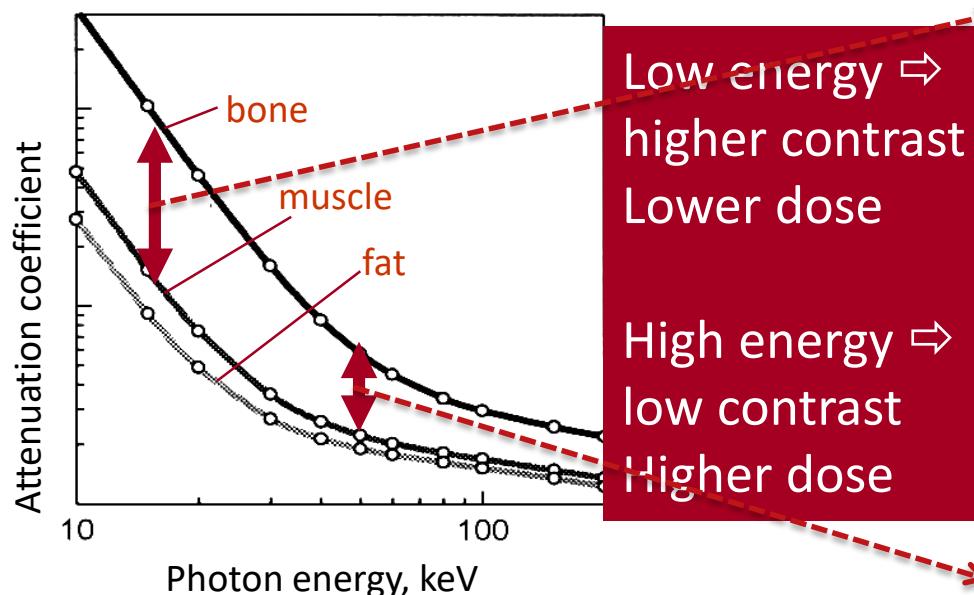
ESD, mGy	U, kV	FSC	Grid factor
0,18	80-100	800	7
0,25	100	400	7
0,35	65-80	800	7
0,44	70-80	400	7
1,04	100	100	7
1,92	70-85	100	7
1,94	85	100	7
2,38	70-75	100	15

13 fold!

Radiography

Image contrast and patient dose

- Absorption depends on photon energy
- The image contrast depends on the energy



Radiography

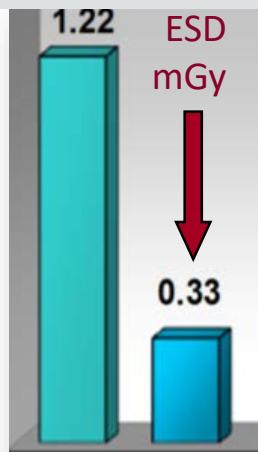
Example: Chest radiography with old radiography system

Chest wall; Bucky $r=7$; $f = 150$ cm; filtration 3 mm Al

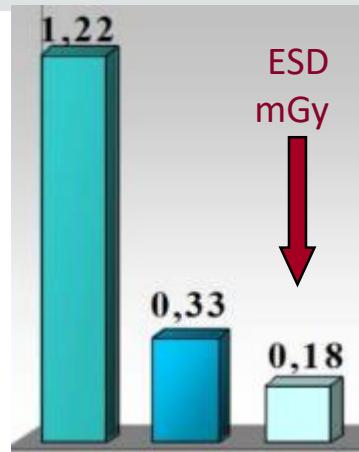
$U = 70 - 80$ kV; speed class 100



Increased speed
class 100 → 400



Increased kVp and filtration
 70 kV → 100 kV,
1 mm Al + 0.1 mm Cu



Radiography

Added filtration

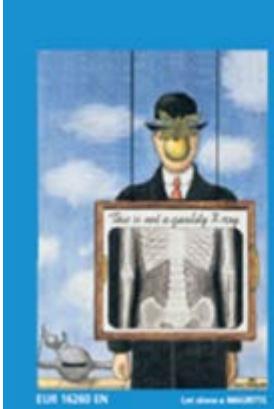
Added filtration	0 mm Al	3 mm Al
Examination	Mean ESD (μ Gy)	Reduction
Abdomen AP 10 months (62 kVp)	200	30 %
Chest AP 10 months (55 kVp)	64	40 %
Pelvis AP 4 months (50 kVp)	94	51 %



From: Mooney and Thomas : Dose reduction in a paediatric X-ray department following optimization of radiographic technique, BJR (77) 1998:852-860

Radiography

Good radiographic technique



3. EXAMPLE OF GOOD RADIOGRAPHIC TECHNIQUE

- | | |
|----------------------------------|---|
| 3.1. Radiographic device: | vertical stand with stationary or moving grid |
| 3.2. Nominal focal spot value: | ≤ 1.3 |
| 3.3. Total filtration: | ≥ 3.0 mm Al equivalent |
| 3.4. Anti-scatter grid: | $r = 10$; 40/cm |
| 3.5. Screen film system: | nominal speed class 400 |
| 3.6. FFD: | 180 (140-200) cm |
| 3.7. Radiographic voltage: | 125 kV |
| 3.8. Automatic exposure control: | chamber selected — right lateral |
| 3.9. Exposure time: | < 20 ms |
| 3.10. Protective shielding: | standard protection |

Radiography

Good radiographic technique

EUROPEAN COMMISSION

**EUROPEAN GUIDELINES
ON QUALITY CRITERIA
FOR DIAGNOSTIC
RADIOGRAPHIC IMAGES
IN PAEDIATRICS**



EUR R(2011)1

PA/AP PROJECTION
(Beyond the newborn period)

For co-operative patients PA projection;
AP projection for non-co-operative patients.

3 EXAMPLE OF GOOD RADIOGRAPHIC TECHNIQUE

- 3.0. Patient position : upright, supine position possible
- 3.1. Padiographic device : table or vertical stand, depending on age
- 3.2. Nominal focal spot value : 0.6 (≤ 1.3)
- 3.3. Additional filtration : up to 1 mm Al + 0.1 or 0.2 mm Cu (or equivalent)
- 3.4. Anti-scatter grid: $r = 8$; 40/cm : only for special indications and in adolescents
- 3.5. Screen film system : nominal speed class 400 - 800
3.6 FFD 100 - 150 cm
- 3.7. Radiographic voltage : 60 - 80 kV (100 - 150 kV with grid for older children)
- 3.8. Automatic exposure control : chamber selected - lateral; preferably none in infants and young children
- 3.9. Exposure time : <10 ms
- 3.10. Protective shielding : lead-rubber coverage of the abdomen in the immediate proximity of the beam edge

Designing the dose audit to obtain the DRL

1. Select examination
2. Select patient data
3. Ensure weight is within required range
4. Record the dose index value
5. Review and clean the data
6. Calculate the median value of dose index



Optimising

- Know the Technology
- Look to implement good radiographic practice

**Establishing DRLs gives a benchmark in
order to optimise equipment and practice**



Establishing and Using DRLs for optimisation in Radiography

Sue Edyvean

ICTP-IAEA Workshop on Establishment and Utilization of Diagnostic Reference
Levels in Medical Imaging Imaging (smr3333):
18-22 November 2019 **Trieste, Italy**

Senior Scientific Group Leader
Medical radiation Dosimetry, CRCE
Public Health England
Didcot, Oxon. OX11 0RQ, UK