# Use of effective dose in medical exposure

2<sup>nd</sup> International ICRP Symposium Abu Dhabi . October 2013

Dr. Pedro Ortiz López ICRP Committee 3



# Overview of the use of effective dose in medical exposure

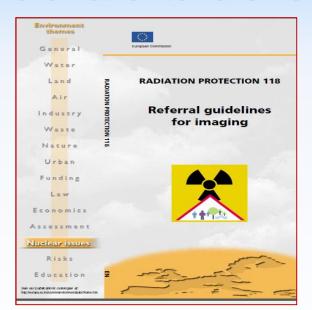
## Effective dose has been useful for protection in imaging medical exposure

- In justification of medical procedures
- In optimization of protection by comparing exposures:
  - Among hospitals, countries
  - For the same type of examination
  - From alternative examinations
- In establishing dose constraints for comforters and volonteers in biomedical research



### In justification

## For the justification of medical exposure, referral criteria exist



CLINICAL PROBLEM	INVESTIGATION {DOSE}	RECOMMENDATION {GRADE}	COMMENT
Orbital lesions  A15	CT (II) or MRI (0)	Specialised investigation (B)	CT provides better anatomical detail, particularly bony structures (e.g. nasolacrimal duct). MR radiation dose to lens (but contraindicated with ferromagnetic FB suspected). Consider US faintra-ocular lesions.
Orbits Metallic FB (before MRI) A16	XR orbits (I)	Indicated (B)	Especially for those who have worked with 1 materials, power tools, etc. Some centres use (see Trauma Section K for acute injury.
Visual disturbances A17	SXR (I)	Not indicated routinely (C)	Plain XRs rarely contributory. Specialists ma CT or MRI.
Epilepsy (adult)	SXR (I)	Not indicated routinely (B)	Evaluation requires specialist expertise. Late seizures should normally be investigated but may be unnecessary if clearly alcohol-related
(for children see Section M) A18	CT (II), MRI (0) or NM (III)	Specialised investigation (B)	Partial/focal seizures may require detailed evif surgery is being considered. Ictal SPECT I likelihood of localising focus. Interictal functimaging also important. Much depends on lowhich will determine combinations of process.

### Typical effective doses from diagnostic medical exposures in the 1990s

Quarter	ROLL CONTROL C	
Water		
Land E	RADIATION PROTECTION 118	
All Se		
Industry &	Referral guidelines	
W g	for imaging	
Natura 2		Diagnostic :
Urban	_	Diagnostic
Punding	A_A	
Law	_	
Economics	10/175	
Attenuent	The second secon	
Minter man		
Risks	- 500	
Education E	- Emerican	
A ALEXANDER OF THE PARTY OF THE	1 - 1 - 1	

Diagnostic procedure	Typical effective dose (mSv)	Equivalent No. of chest x-rays	Approximate equivalent period of natural background radiation (1)
X-ray examinations:			
Limbs and joints (except hip)	< 0.01	<0.5	<1.5 days
Chest (single PA film)	0.02	1	3 days
Skull	0.07	3.5	11 days
Thoracic spine	0.7	35	4 months
Lumbar spine	1.3	65	7 months
Hip	0.3	15	7 weeks





Barium swallow	1.5	75	8 months
Barium meal	3	150	16 months
Barium follow through	3	150	16 months
Barium enema	7	350	3.2 years
CT head	2.3	115	1 year
CT chest	8	400	3.6 years
CT abdomen or pelvis	10	500	4.5 years
Radionuclide studies:			
Lung ventilation (Xe-133)	0.3	15	7 weeks
Lung perfusion (Tc-99m)	1	50	6 months
Kidney (Tc-99m)	1	50	6 months
Thyroid (Tc-99m)	1	50	6 months
Bone (Tc-99m)	4	200	1.8 years
Dynamic cardiac (Tc-99m)	6	300	2.7 years
PET head (F-18 FDG)	5	250	2.3 years



# Broad categories



## TABLE Classification of the typical effective doses of ionising radiation from common imaging procedures

Class	Typical effective Dose (mSv)	Examples
0	0	US, MRI
I	<1	CXR, limb XR, pelvis XR
Ш*	1–5	IVU, lumbar spine XR, NM (e.g. skeletal scintigram), CT head & neck
III	5–10	CT chest and abdomen, NM (e.g. cardiac)
IV	>10	Some NM studies (e.g. PET)

<sup>\*</sup> The average annual background dose in most parts of Europe falls in Band II.

## Optimization

### Comparisons of effective dose

- The same type of examination with the same technique, but taken in different rooms, hospitals, or countries
- The same type of examination obtained with different techniques or projections
- Different types of examinations (for example a nuclear medicine with a CT examination)



## Optimization situation 1: same type of examination, same exposure technique

- The dose distribution inside the body is similar
- There is no need to calculate organ doses nor effective dose to compare
- Comparisons can be made directy in terms of measurable quantities. Examples:
  - entrance surface air kerma,
  - air kerma-area product,
  - administered activity of radiopharmaceutical



## Optimization situation 2: comparing different techniques

- Different exposure factors, different projections, different radiopharmaceuticals
- The dose distribution is different
- In this situation effective dose is appropriate for comparison



### Professional guidelines relating to optimization

### **European Association of Nuclear Medicine**

Eur J Nucl Med Mol Imaging DOI 10.1007/s00259-007-0694-9

**GUIDELINES** 

EANM/ESC guidelines for radionuclide imaging of cardiac function

B. Hesse • T. B. Lindhardt • W. Acamna •





400,000 particles will result in obstruction of only a very small fraction of pulmonary vessels. A reduction in the number of particles administered to between 100,000 and 200,000 is recommended in patients with known pulmonary hypertension, right to left heart shunt or after a single

suggested. Perfusion-only scans should be performed on day 1, using a reduced dose of <sup>99m</sup>Tc-MAA. In most patients PE can be excluded on the basis of a normal perfusion pattern. When the perfusion pattern is abnormal but not diagnostic of PE, subcutaneous low molecular

Table 1 Data on radiation exposure in adults

Reference	Radiopharmaceutical	Administered activity (MBq)	Critical organ, dose (mGy/MBq)	Effective dose (mSv/MBq)
[71]	<sup>99m</sup> Tc-MAA	40–120	Lungs, 0.067	0.017
[72]	<sup>99m</sup> Tc-DTPA	20-30	Bladder, 0.047	0.007
[73]	Technegas	20-30	Lungs, 0.11	0.015
[74]	<sup>81m</sup> Kr	40-400	Lungs, 0.0068	0.0007

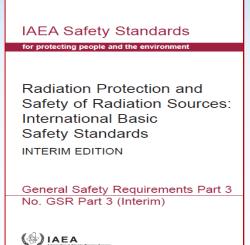
Critical organ, dose (mGy/MBq)	Effective dose (mSv/MBq)
Lungs, 0.067	0.017
Bladder, 0.047	0.007
Lungs, 0.11	0.015
Lungs, 0.0068	0.0007



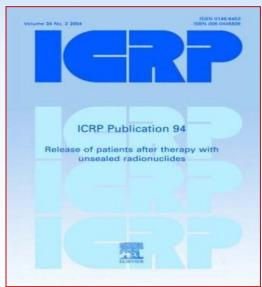
# Dose constraints for comforters and volunteers

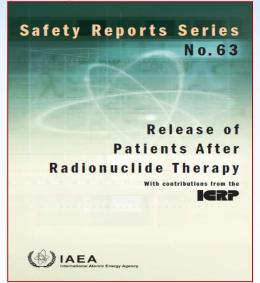
### Dose constraints to comforters

International Safety
 Standards establish
 requirements on
 constraints for comforters
 and volonteers in
 biomedical research,
 when the exposed
 individual does not
 directly benefit from the
 exposure



Guidance based on efective dose is used for this purpose





# Are there other uses of effective dose other than strictly radiation protection?

### SOURCES AND EFFECTS OF IONIZING RADIATION

United Nations Scientific Committee on the Effects of Atomic Radiation

> UNSCEAR 2008 Report to the General Assembly with Scientific Annexes

> > VOLUME I



## Effective dose for summarizing uses of radiation and contributions to total exposure

- Average effective dose per procedure
- Collective effective dose for a given procedure or practice or for the whole x-ray diagnostic or nuclear medicine
- Per caput doses, by dividing collective effective dose by the population



# Effective dose, collective effective dose UNSCEAR 2000

Table 29
Some reported annual individual and collective effective doses from diagnostic medical x-ray examinations <sup>a</sup>
Data from UNSCEAR Survey of Medical Radiation Usage and Exposures unless otherwise indicated

_	Effective do	ose (mSv)	Collective effective dose	
Country	Per examination	Per caput	(man Sv)	Ref.
		Health-care level I		
Australia	1.3	0.8	13 000	[W34]
Bulgaria	1.28	0.75	6 400	- 1
Canada	1.05	0.94	26 200	[A15]
China, Taiwan Province	0.43	0.23	4 700	[L23]
Denmark	0.7	0.36	1 820	- 1
Finland	0.63	0.45	2 270	-
France	-	1.0	57 660	[S50]
Germany	1.5	1.9	153 360	
Netherlands	1.0	0.6	9 000	-
Poland	1.2	0.8	32 300	-
Portugal	0.83	0.71	7 000	[F11]
Romania	1.35	0.61	13 800	
Russian Federation	0.7	0.9	128 000	-
Sweden	1.2	0.68	6 000	-
Ukraine	0.83	0.50	26 250	[K18]



## Example 2: contribution to collective (effective) dose (UNSCEAR 2000)

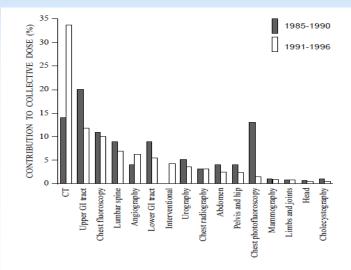


Figure V. Percentage contributions by examination type to global collective dose from medical x-ray examinations: comparison of data for 1955-1990 and 1991-1996.

### Per caput dose (collective effective dose divided by the total population, UNSCEAR 2000)

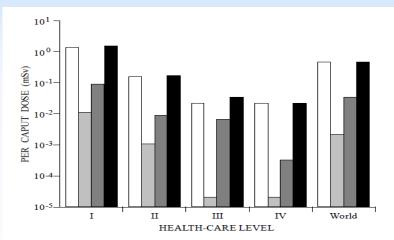


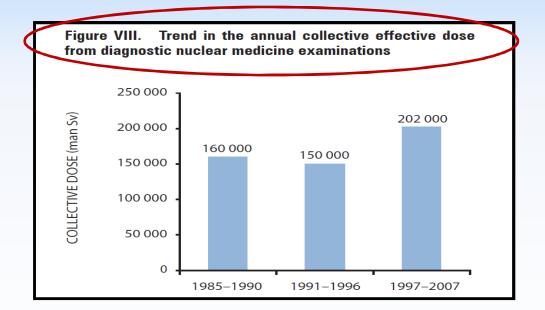
Figure IX. Estimated global annual per caput doses from medical diagnostic radiological procedures (1991-1996). The four columns in each group represent medical x rays, dental x rays, nuclear medicine (diagnosis), and all diagnostic practices, respectively.

### **UNSCEAR 2008**

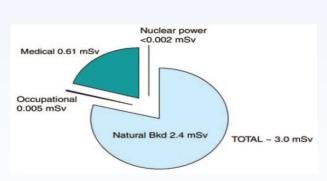
Table 1. Estimated annual per caput dose and annual effective dose to the world population from diagnostic medical and dental radiological examinations (1997–2007)

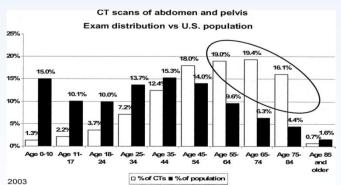
Health-care level	Population (millions)	Annual per caput dose (mSv)		Annual collective effective dose (man Sv.	
		Madi	Dental	Medical	Dental
I	040	1.91	0.006 4	2 900 000	9 900
Anr	nual collec	tive eff	fective di	ose (man	SW
7 1171	raar oonoc	ACIVO OII		ooo man	0.,
Medical				Denta	3/
20	000 000			0.00	n
/ 5	14.74.7 4.74.74.7		I		

### **UNSCEAR 2008**



# Comparison of collective effective doses are often for different age and sex distributions



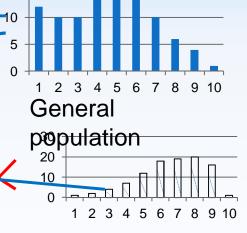




Effective dose involves tissue weighting factors,  $w_T$ , and detriment risk coefficients, r

$$E = \sum_{T} \sum_{R} w_{T} w_{R} D_{T,R}$$
• Detriment =  $PE$ 

 Coefficients w<sub>T</sub> and r were derived for populations and for the general population



Example of patient population



### Question (caveat):

- Effective dose is used with some flexibility for populations that are different from those for which  $w_T$  and effective dose are derived
- In the context of
  - summarizing uses of radiation in medicine and
  - contribution to the population exposure,
  - not to calculate risk



Effective dose should be expressed with no more than two significant figures



#### UNSCEAR 2000 Report (page 296-297)

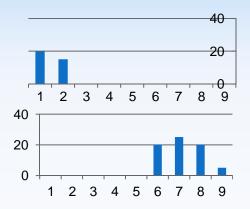


- 1. "effective dose should not be used directly for estimating detriment from medical exposure (individuals or populations) ... by application of the nominal fatality probability coefficients. Such assessments would be inappropriate and serve no purpose in view of the uncertainties arising from potential demographic differences (in terms of health status, age and sex), between particular population of patients and those from general populations for whom ICRP derived the risk coefficients ... effective dose could broadly underestimate the detriment from diagnostic exposures of young patients by a factor of 2 and, conversely, could overestimate the detriment from old patients by a factor of at least 5.
- 2. "The analysis of radiation risk from diagnostic medical exposures requires detailed knowledge of organ doses and the age and sex of patients...."
- 3. "It is possible...to use effective dose and even collective dose for medical diagnostic exposure as long as this is done only for comparative purposes and for the same or similar patient populations, and it would require additional considerations or significant corrections if we try to use them to compare with other populations."





### Skewed populations,



**Pediatric** 

Old adults

For comparisons for these different population groups, corrections may be needed



# Recap: Effective dose is useful in medical exposure

- For radiation protection:
  - Justification
  - Optimization
  - Dose constraints for comforters and volonteers
- For expressing radiation use and contributions to population exposure
- Butõ



### But, remembering that

 Collective effective dose is not intended as a tool for epidemiological risk assessment, and it is inappropriate to use it in risk projections (ICRP 103).

