# PATIËNTEN DOSIMETRIE VOOR DETECTIE EN FOLLOW-UP VAN SUBOPTIMALE PROCEDURES

VBS-GBS Radioprotectie, Brussel September 21, 2019

N Buls Radiologie

Nico.Buls@uzbrussel.be









RADIO protectie protection

21.09.2019

# FOCUS

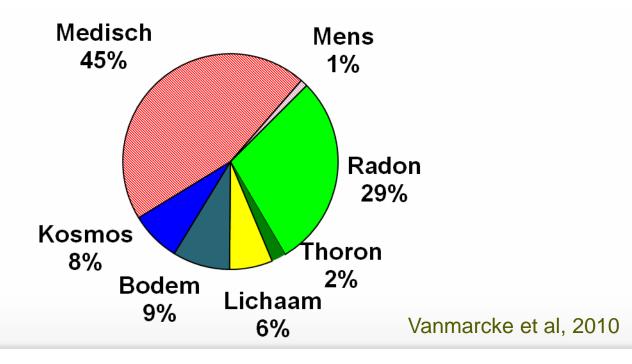
1. Level of patient exposure during medical procedures with x-ray imaging. Are dose values in your practice "acceptable" or are they "too high".

- 2. Specific dosimetric quantities for medical exposure. Which quantities should I consider? Which are available?
- 3. How can I improve quality by dose assessment?





- ~ 2,5 mSv/y natural sources
- ~ 2,0 mSv/y medical imaging of which ~50% from CT







**EUROPEAN COMMISSION** 

#### RADIATION PROTECTION Nº 180

Medical Radiation Exposure of the European Population

Part 1/2

Directorate-General for Energy Directorate D — Nuclear Safety & Fuel Cycle Unit D3 — Radiation Protection 2014

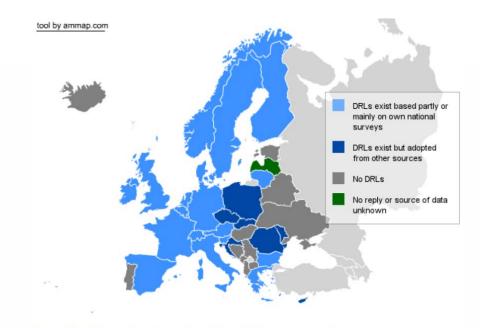
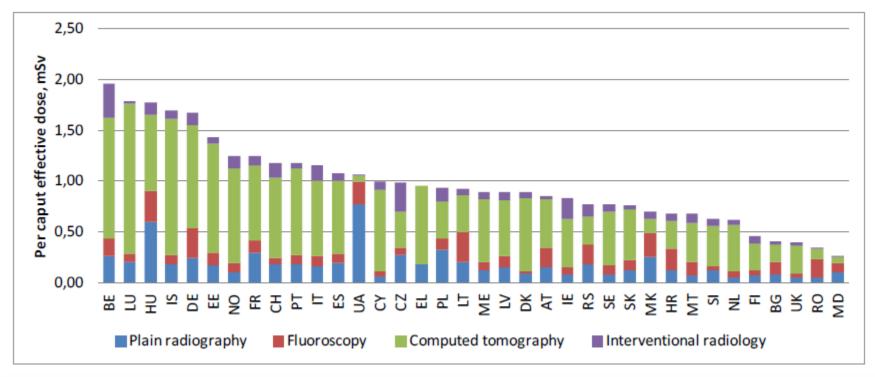


Figure 2.1. Diagnostic reference levels for adult x-ray examinations.



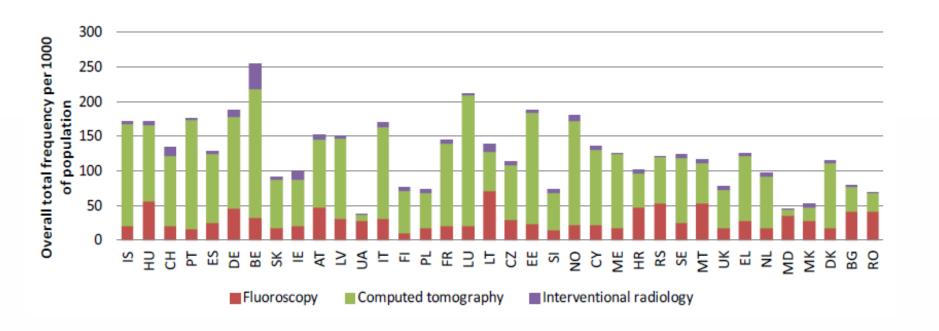




Source Study on European Population Doses from Medical Exposure (Dose Datamed 2, DDM2), 2014







Source Study on European Population Doses from Medical Exposure (Dose Datamed 2, DDM2), 2014





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# Effective Doses in Radiology and Diagnostic Nuclear Medicine:

A Catalog<sup>1</sup>

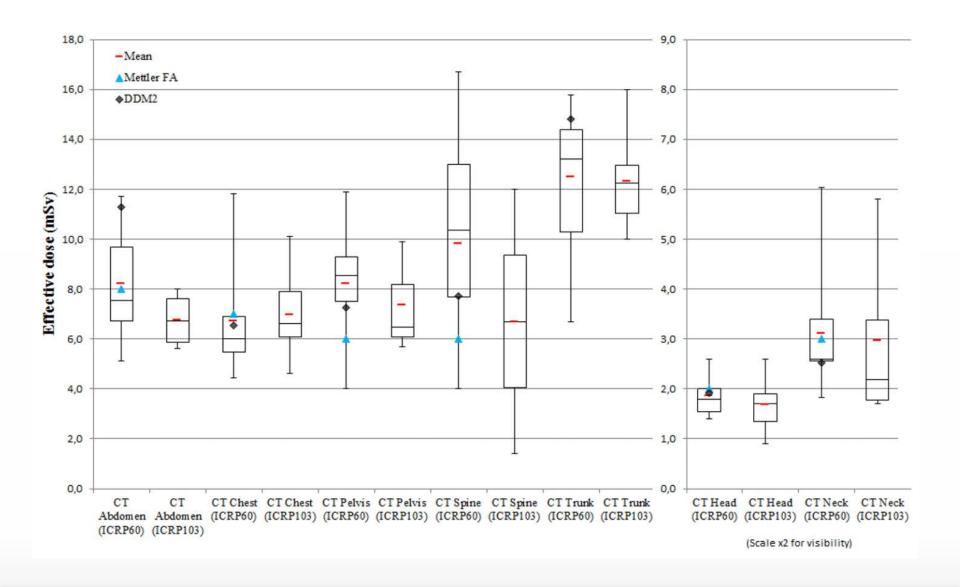
Fred A. Mettler, Jr, MD, MPH Walter Huda, PhD Terry T. Yoshizumi, PhD Mahadevappa Mahesh, MS, PhD

Medical uses of radiation have grown very rapidly over the past decade, and, as of 2007, medical uses represent the largest source of exposure to the U.S. population. Most physicians have difficulty assessing the magnitude of exposure or potential risk. Effective dose provides an approximate indicator of potential detriment from ionizing radiation and should be used as one parameter in evaluating the appropriateness of examinations involving ionizing radiation. The purpose of this review is to provide a compilation





Radiology







# THE USE OF EFFECTIVE DOSE FOR MEDICAL EXPOSURE



The estimation of effective dose can be problematic in medical exposure.

(341) The assessment and interpretation of effective dose from medical exposure of patients is problematic when organs and tissues receive only partial exposure or a very heterogeneous exposure, which is the case especially with diagnostic and interventional procedures.

The inherent uncertainty for organ dose estimation in a **reference** patient is around  $\pm$  40% c. Martin, Br J Radiol (2007)





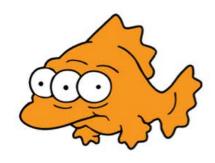
# SIMPSONS GUIDE TO RADIATION DOSIMETRY



Bequerel [Bq] How brightly your Cesium glows



Gray [Gy]
How brightly
Cesium will make
you glow



Sieverts [Sv]
How many extra
eyes will you have
after glowing?





# RADIATION DOSE QUANTITIES AND UNITS

#### Physical quantities

Describe physical effect
Can be measured directly
Fluence, Kerma (gray), Absorbed dose, *D* (gray)

#### Radiation Protection quantities

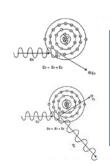
To consider health effects, to set legal dose limits Can not be measured Organ equivalent dose,  $H_T$  (sievert), Effective dose, E (sievert)

#### Operational quantities

Measurable for compliance with dose limits Area monitoring and individual monitoring  $H_p(d)$ , d = 10 mm

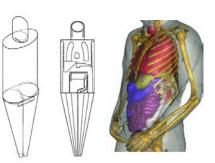






#### Primary physical quantities

Kerma (gray) Absorbed dose, *D* (gray)



Calculated using simple phantoms (sphere or slab) validated by measurements

#### Operational quantities

Ambient dose equivalent (Sv) Personal dose equivalent,  $H_p(10)$  (Sv)

Related by calibration and calculation





Monitored quantities
Instrument responses

#### By Simulation

- Human model
- Radiation  $w_r$  and tissue  $w_t$  weighting factors

#### Protection quantities

Organ dose, (Sv) Effective dose, (Sv)

Relate to

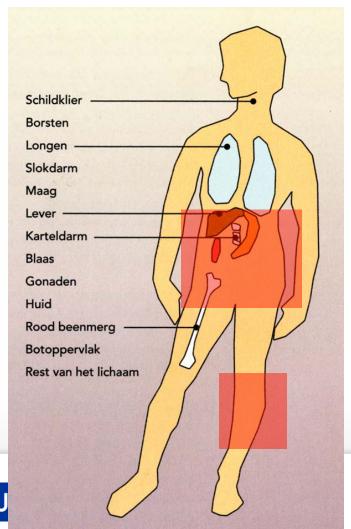
Radiation health risk

Cancer risk, tissue effects, etc





# EFFECTIVE DOSE (SIEVERT)



Effective dose is the sum of all organ doses corrected for differences in organ sensitivity (tissue weight factors).

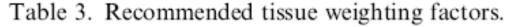
$$E = \sum w_T \times H_{T,R} \quad [Sv]$$

Related to the biological effect

Depends on exposed region

# TISSUE WEIGHTING FACTORS

# ICRP report 103 recommendations (2007)

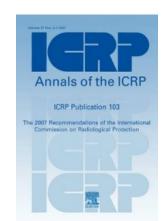


Tissue	$w_{\mathrm{T}}$	$\sum w_{\mathrm{T}}$
Bone-marrow (red), Colon, Lung, Stomach,	0.12	0.72
Breast, Remainder tissues*		
Gonads	0.08	0.08
Bladder, Oesophagus, Liver, Thyroid	0.04	0.16
Bone surface, Brain, Salivary glands, Skin	0.01	0.04
	Total	1.00

<sup>\*</sup> Remainder tissues: Adrenals, Extrathoracic (ET) region, Gall bladder, Heart, Kidneys, Lymphatic nodes, Muscle, Oral mucosa, Pancreas, Prostate (♂), Small intestine, Spleen, Thymus, Uterus/cervix (♀).







# TISSUE WEIGHTING FACTORS

The knowledge of radiation effects is constantly evolving epidemiological evidence, exposed populations radiobiological evidence

Scientific progress is not fully reflected in present legislation and recommendations

Tissue weighting coefficients depend on our current knowledge of radiobiology





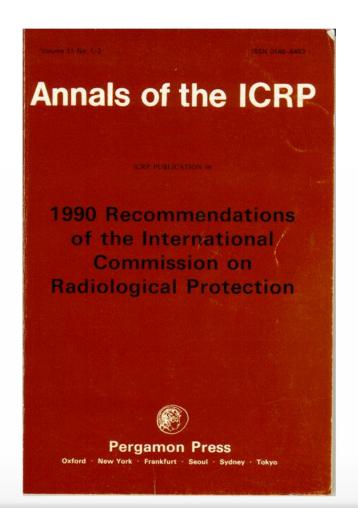
# REGULATORY FRAMEWORK

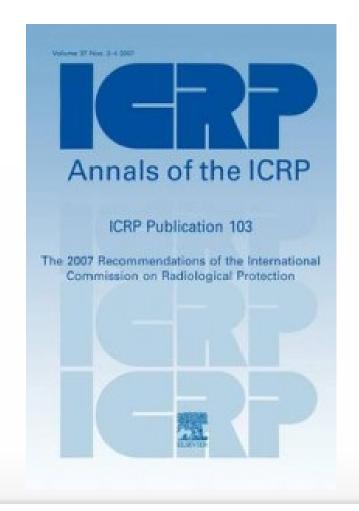






# REGULATORY FRAMEWORK

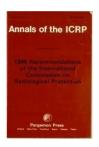








# **EVOLUTION TISSUE FACTORS**





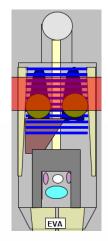
	ICRP 60, 1990	ICRP 103, 2007
Breast	0.05	0.12
Gonads	0.20	0.08
RBM, lung, colon, stomach	0.12	0.12



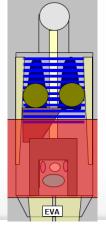


# CONSEQUENCE...





Examples	ICRP 60, 1990	ICRP 103, 2007
Cardiac CT 120 kVp, 16cm, DLP = 200 Gycm	4.1 mSV	5.4 mSv
Abdomen CT, 1 sequence 120 kVp, 50cm, DLP = 800 Gycm	15 mSv	12 mSv

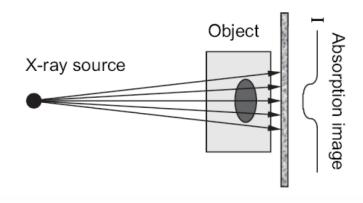






# EFFECTIVE DOSE FROM X-RAY'S?





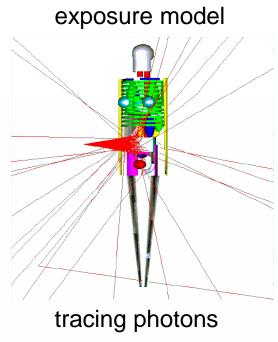
$$I = \int_{0}^{E \max} I_0(E) e^{-\int_{0}^{d} \mu(E) \cdot ds} dE$$



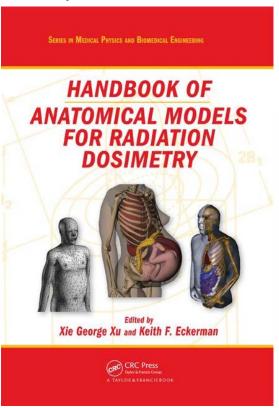


#### FROM GRAY TO SIEVERT IN MEDICAL IMAGING





### patient model

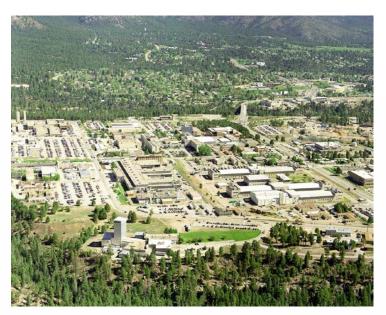






#### COMPUTATIONAL PHYSICS: TRACING PHOTONS BY SIMULATION













#### COMPUTATIONAL PHYSICS: TRACING PHOTONS BY SIMULATION



Founded during World War II as a secret, centralized facility to coordinate the scientific research of the Manhattan Project

#### J. Robert Oppenheimer



First director (1943-1945)

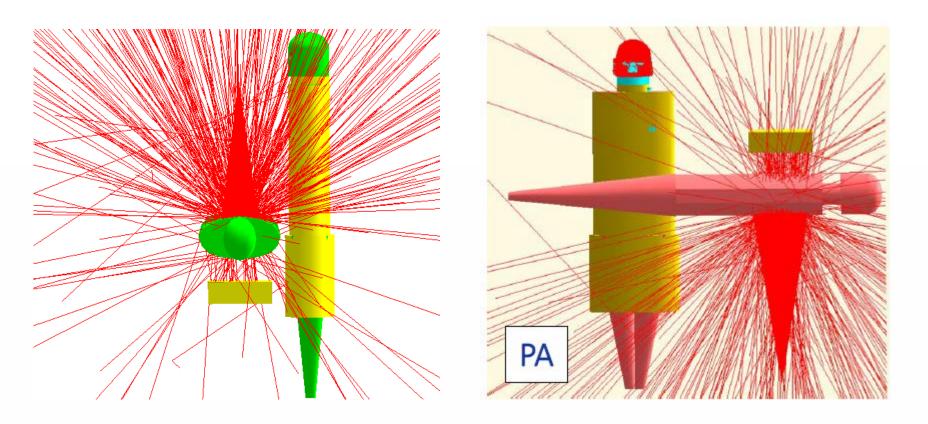
Today one of the largest science and technology institutions in the world. Multidisciplinary research in fields such as

- national security
- space exploration
- nuclear fusion
- renewable energy
- medicine
- nanotechnology
- supercomputing





#### TRACING PHOTONS BY MONTE CARLO SIMULATION MODELS

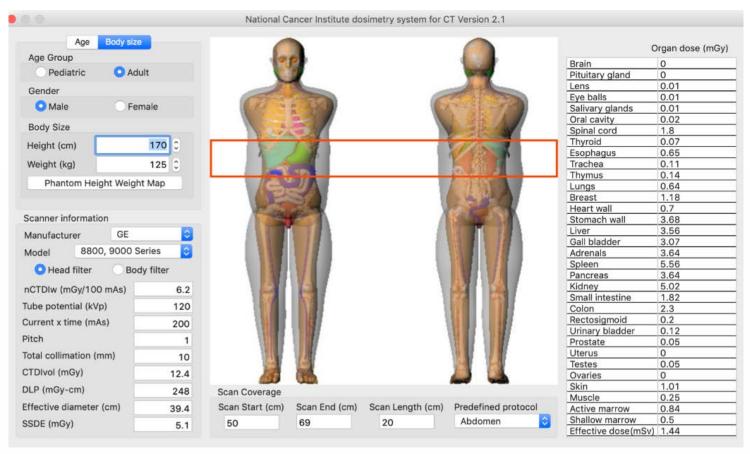


Photon transport simulations with Monte Carlo N-Particle Transport Code





#### FROM GRAY TO SIEVERT IN MEDICAL IMAGING - EXAMPLE CT



https://ncidose.cancer.gov/#ncict





#### GENERIC CONVERSION FATORS – EXAMPLE CT

#### DLP to E conversion coefficients

TABLE 3 Normalised values of effective dose per dose-length product (DLP) over various body regions and (standard) patient ages (Shrimpton, 2004)

Region of body	Effective dose per DLP (mSv (mGy cm) <sup>-1</sup> ) by age					
	0 y oldª	1 y oldª	5 y oldª	10 y oldª	Adult <sup>b</sup>	
Head & neck	0.013	0.0085	0.0057	0.0042	0.0031	
Head	0.011	0.0067	0.0040	0.0032	0.0021	
Neck	0.017	0.012	0.011	0.0079	0.0059	
Chest	0.039	0.026	0.018	0.013	0.014	
Abdomen & pelvis	0.049	0.030	0.020	0.015	0.015	
Trunk	0.044	0.028	0.019	0.014	0.015	

#### Notes:





<sup>&</sup>lt;sup>a</sup>All data normalised to CTDI<sub>w</sub> measured in the 16 cm diameter CT dosimetry phantom.

<sup>&</sup>lt;sup>b</sup>Data for the head & neck regions normalised to CTDI<sub>w</sub> in the 16 cm diameter CT dosimetry phantom; data for other regions normalised to CTDI<sub>w</sub> in the 32 cm diameter CT dosimetry phantom.

#### EFFECTIVE DOSE – EXAMPLE CT

#### AJ Einstein et al, Radiology 2010

Method Used	Effective Dose Derived from	Helical	Helical ESTCM	Prospective Helical	Volume with Standard Exposure Time	Volume with Optimized Exposure Time	Volume 100 kVp: Optimized Exposure Time
ICRP publication 103 <sup>†</sup>	Organ doses	35.4	22.3	9.3	8.2	5.8	4.4
ICRP publication 60	Organ doses	26.5	16.6	7.0	5.9	4.1	3.1
2000 European guidelines	$DLP \times k$ factor	20.4	14.0	5.9	4.8	3.2	2.2
2004 European quidelines	$DLP \times k$ factor	16.8	11.6	4.9	4.0	2.7	1.8
DLP (mGy·cm)		1201.3	826.2	348.9	284.7	189.8	128.6
Dose reduction from helical (ICRP publication 103) (%)			37.0	73.7	76.8	83.6	91.2

# Same scan, difference in effective dose x 2





# PATIENT DOSE REGISTRATION

Okay, we have to be careful when we use **effective dose** in medical exposure



Yes, we have to consider other, practical metrics that can be easily **measured** 





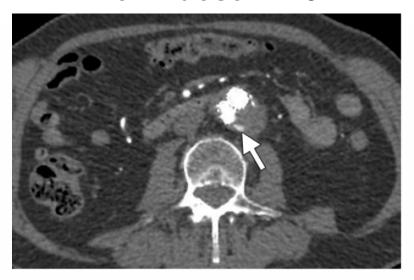
# PRACTICAL DOSIMETRY FOR

# **Optimisation and good practice (ALARA)**

"Standard" dose



"Low" dose ~ ÷ 5



Effect of technical parameters (kVp, mA, s, pulse time, pitch, collimation, etc) on dose?





# PRACTICAL QUANTITIES FOR MEDICAL EXPOSURE

We require PRACTICAL radiation dose quantities for patient dosimetry from medical exposures

**Immediate feedback**: indicated before (estimation), during and immediately after the examination.

dose indicators characterizing radiation exposure in imaging for the purposes of **comparison of practice**.

these quantities are **not patient doses** (directly reflecting **risk to individuals**)

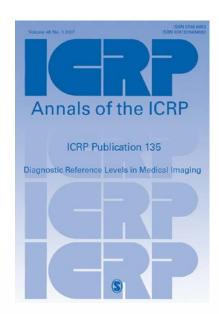




# DIAGNOSTIC REFERENCE LEVELS (DRL)

Diagnostic reference level (DRL)

A diagnostic reference level is a form of investigation level used as a tool to aid in optimisation of protection in the medical exposure of patients for diagnostic and interventional procedures. It is used in medical imaging with ionising radiation to indicate whether, in routine conditions, the amount of radiation used for a specified procedure is unusually high or low for that procedure. For



#### DRL quantity

A commonly and easily measured or determined radiation metric (e.g.  $K_{a,e}$ ,  $K_{a,i}$ ,  $CTDI_{vol}$ , DLP,  $P_{KA}$ ,  $K_{a,r}$ ,  $D_G$ , administered activity) that assesses the amount of ionising radiation used to perform a medical imaging task. The quantity or quantities selected are those that are readily available for each type of medical imaging modality and medical imaging task. Suitable quantities





# PRACTICAL DOSIMETRY FOR

# Dose archiving, reporting and benchmaring



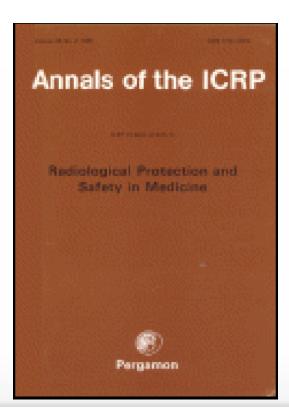
#### **Diagnostic Reference Levels (DRL)**





# PRACTICAL DOSIMETRY FOR

# Dose archiving, reporting and benchmaring





Council directive 97/43/Euratom of 30 juin 1997 on health protection of individuals against the dangers of ionizing radiation in relation to medical exposure and repealing directive 94/466/Euratom.

Art 4: Member States promote the establishment and the use of diagnostic reference levels for radiodiagnostic examinations

ICRP report 73 (1996)





# LEGAL BASIS



FANC decree: 28.09.2011 (BS 11.10.2011)

Conduct periodical studies to determine Diagnostic Reference Levels (DRL) for standard examinations



CT: yearly



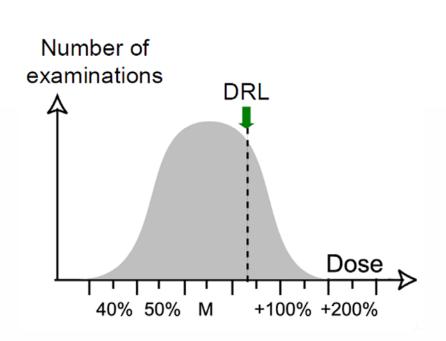
RX en IR: 3-yearly

In each center, the average dose should be compared to the latest DRL for optimization purposes





# OPTIMISING CLINICAL PRACTICE BY DRL





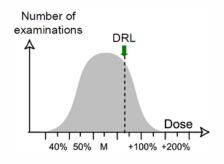
75 percentile of dose distribution

**Iterative process** 





# OPTIMISING CLINICAL PRACTICE BY DRL



#### · DRN's bij volwassenen

VOLWASSENEN	CTDI <sub>vol</sub> (mGy)		DLP (mGy.cm)			
VOLWASSENEN	Enkelv	oudig onderzoek	Enkelvoudig onderzoek		Volledig onderzoek	
Onderzoek	P25	DRN (P75)	P25	DRN (P75)	P25	DRN (P75)
Abdomen	7	10	320	490	350	570
Angio CT van de thorax	5	11	165	300	180	330
Hart (CCTA)	10	25	140	290	190	480
Colon	3*	4*	150*	200*	230	460
Cervicale wervelzuil	14	25	230	450	-	-
Lumbale wervelzuil	18	26	380	600	-	-
Schedel (hersenen)	35	50	640	900	-	-
Sinussen	2,5	6	35	80	-	-
Thorax	5	8	170	260	-	-
Thorax - abdomen	5	8,5	320	550	480	800





## RADIATION DOSE QUANTITIES MEDICAL EXPOSURE

## **Depends on modality**





air kerma at reference point (mGy) dose-area product DAP (mGy.cm²)



the CT dose index *CTDIvol (mGy)* dose length product *DLP (mGy.cm)* 



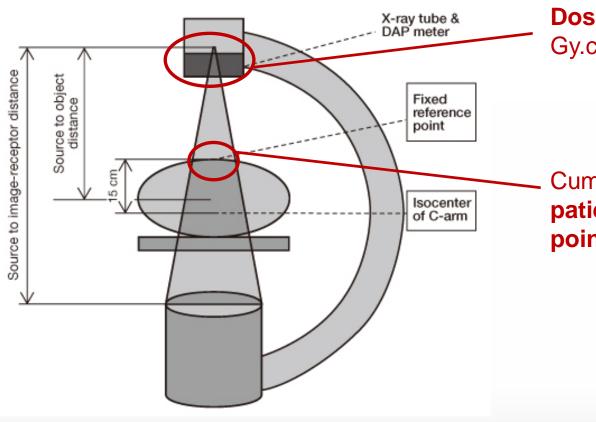
entrance skin dose *ESD* (*mGy*)
Average Glandular Dose *AGD* (*mGy*)





## RADIOGRAPHY AND FLUOROSCOPY

## Two operational units



Dose Area Product (DAP) Gy.cm<sup>2</sup>

Cumulative dose (CD) in **patient entrance reference point** in mGy

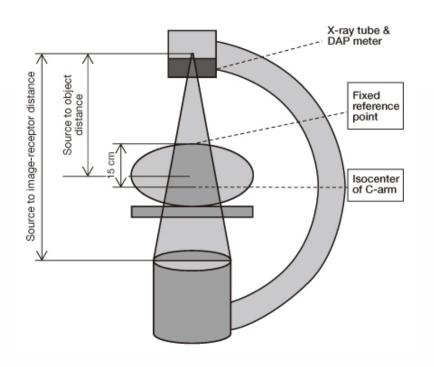
represent the air kerma incident on the patient's skin surface



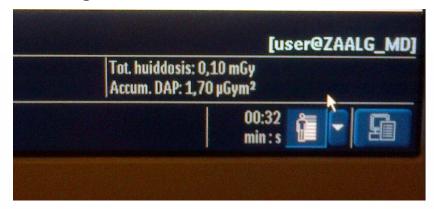


## RADIOGRAPHY AND FLUOROSCOPY

## Two operational units



#### **During examination**



## **During fluoroscopy**

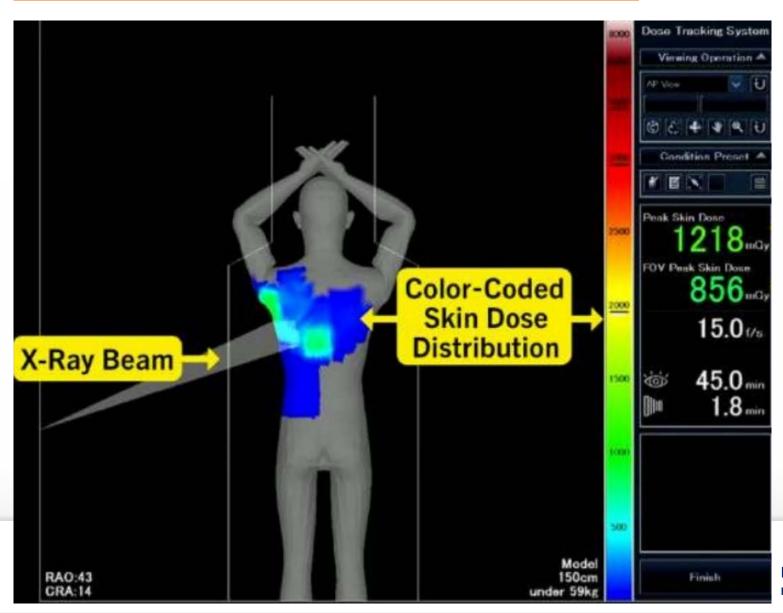






# Real time skin dose mapping

## RADIOGRAPHY AND FLUOROSCOPY



Example from medical canon

## COMPUTED TOMOGRAPHY

## Two operational units

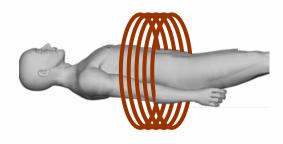
#### Dose from one tube rotation

- weighted CT dose index (CTDI<sub>w</sub>)
- volume weighted CT dose index (CTDI<sub>vol</sub>)



### Dose from one sequence

- dose-length product (DLP)

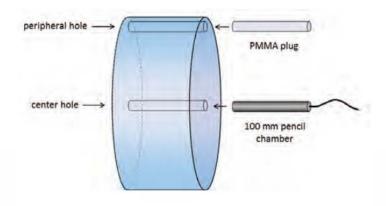






## COMPUTED TOMOGRAPHY

CT systems indicate CTDI and DLP doses based on measurements in **standardized** PMMA phantoms



Does not directly reflect dose in patient (patient size, ...)



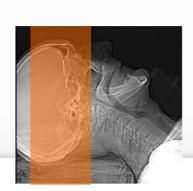




## COMPUTED TOMOGRAPHY







**Dose Information** 

DLP

mGy-cm

353.31

Dose

Eff. %

87.40

Phantom

cm

Head 16

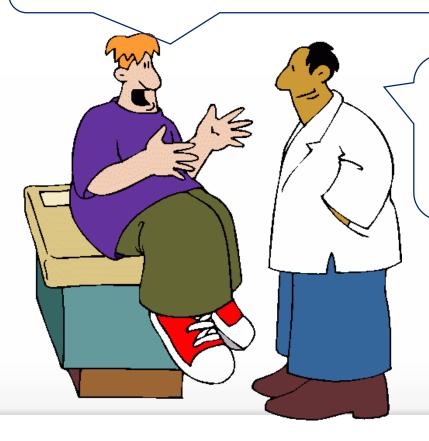
**CTDIvol** 





## PATIENT DOSE REGISTRATION

Now, I understand which dosimetric quantities I have to measure



Okay, let's take a look on how we can record and archive them





## REGISTRATION OF PATIENT DOSES

nde	icatie zaal: tor: 1 Isatste DA rzoek:	R.K	Belli	č					Gener Gener	C-50	* enkel indien  ** facultatief  *** duid de ee  ****gecorrigee  \$\$D (Skin So	nheid aan eerde dosis	geef eenhe	id op)	
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1	alalus	V	-3.4.	97	10,2					1	7.18				
2_	52/01/12	_ <u>V</u> _	43.	97	9,8						62.8				
4	arl dee	n	72	-37	15					1	1228				
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6	al-du	H	78	97	16,6			$ \overline{}$	V	<u>ا</u>	N28				
7	eglodis	H	48	97	194					1	217				
8	Boiles	W	37	37	8	<b>—</b>				1	563				11:
9	11	V	52	37	12 4	_				1	343				
10		77	63	24	108	/				1	7-28				
en o	om gevoeg ndosimetri	jd te wo	orden bij	het bes	luit van	het Fed	eraal Ag	entsch	ap voor n	ucleaire o	ontrole van 2	28 septen	nber 2011	betreffende	
sel,	28 septen	nber 20	11.												-
	teur-gene	la es													







#### BIJLAGE 5. Registratieformulier voor patiëntendosisgegevens - CT-onderzoeken bij minimum 20 patiënten per procedure of gedurende 3 maanden (jaarlijkse dosisstudies)

Centr	um:			UZ Brus	sel						]		
Merk	scanner:			GE Heal	lthcare						1		
Type:	scanner:			Revolut	ion CT						1		
Aanta	l slices:			256 slice	es - 16 cm						,		
Iterati	eve reconstr	uctie softv	Jare:	ASIR-V						İ			
	rzoek:			Hart (0	CTAI					ł			
					,					J			
C	ctpersoon:		Marin.	Gort Vo	Gompel		Voornaam:			1	gort uano	ompol@u	ızbrussel.be
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Fello	ue:		van			ube Current Mo			* duid gebruik	. OTDI.	]		
Nr	Datum	Pati	ent	k¥p	TCM	lteratieve	Low	Contrast-	CTDI_ 10	DLP	Aantal	nitialer	Opmerking(en)
	dd/mm/jiji			(kV)	(JA	econstructi	dose	injectie	сты	(mGu.cm)			Opmerking(en)
		(M of V)	0	(,	of NEE)	(JA of NEE)	(JA of NEE)	(JA of NEE)	(mGy)	,,	1		
1	19-10-2018	М	43,06	100	JA	JA	JA	JA	16,97	364,82	1		
2	19-10-2018	М	64,1	100	JA	JA	JA	JA	7,34	172,48	1		
3	19-10-2018	М	51,94	100	JA	JA	JA	JA	14,09	366,38	1		
4	10/10/2018	٧	69,05	100	JA	JA	JA	JA	5,05	100,92	1		
- 5	9/11/2018	M	52,55	100	JA	JA	JA	JA	6,83	160,44	1		
- 6	9/07/2018	V V	74,4	120	JA JA	JA	JA	JA	24,96	411,80	1		
7 8	9/07/2018	M	77,82 76,48	120	JA JA	JA JA	JA JA	JA JA	2,37 4,32	37,94 60.45	1		
9	9/05/2018	M	71,4	120	JA	JA	JA	JA	7,14	149,96	1		
10	9/04/2018	M	78,47	100	JA	JA	JA	JA	10,31	242,28	1		
- 11	30-08-2018	M	44,41	120	JA	JA	JA	JA	11,01	242,20	1		
12	29-08-2018	М	57,55	120	JA	JA	JA	JA	21,79	414,08	1		
13	28-08-2018	٧	80,98	100	JA	JA	JA	JA	7,46	149,26	1		
14	28-08-2018	٧	74,46	120	JA	JA	JA	JA	10,49	167,88	1		
15	27-08-2018	М	71,6	120	JA	JA	JA	JA	29,43	474,49	1		
16	24-08-2018	М	62	120	JA	JA	JA	JA		71,28	1		
17	23-08-2018	M	56,86	100	JA	JA	JA	JA		225,85	2		
19	23-08-2018	M V	56,86 83,39	100	JA JA	JA JA	JA J	$\Box$	3,67	381,64 173,41	1		
20	14-08-2018	M	38,88	100	JA	JA JA	- A	.10	5,96	128,13	1		
21	13-08-2018	V V	63,98	120	JA	JA	JA		2,10	33,62	1		
22	13-08-2018	м	72,41	100	JA	JA	JA	JA	15,93	374,38	1		
23	8/08/2018	٧	62,1	120	JA	JA	JA	JA	3,13	43,79	2		
24	8/08/2018	٧	62,1	100	JA	JA	JA	JA	17,21	404,54			
25	8/06/2018	М	63,76	100	JA	JA	JA	JA	13,32	306,25	1		
26	8/03/2018	М	19,06	100	JA	JA	JA	JA	5,15	121,08	1		
27	8/03/2018	٧	64,92	100	JA	JA	JA	JA	7,59	163,14	1		
28	8/02/2018	V	80,27	100	JA	JA	JA	JA	12,86	257,11	1		
29	8/02/2018	٧	40,2	120	JA	JA	JA	JA	3,71	51,97	1		
30	8/02/2018	M	72,15	100	JA	JA	JA	JA.	9,87	222,09	1	<b>—</b>	
31	7/09/2018	M M	59,38 57,98	120	JA JA	JA JA	JA JA	JA JA	11,70	275,03 290,61	1		
33	26-06-2018	M	67,83	100	JA	JA JA	JA	JA.	10,42	244,85	1		
34	13-06-2018	M	80,18	100	JA	JA	JA	V-1	9,70	213,30	1		
			1.7/110						-,,,,,				

9,57

4,73 101,78

12,92 297,18

10,51

3,69 68,29

10,84 232,98

5,78 92,54

224,98

168,14

36 6/06/2018

37 6/05/2018

38 6/04/2018

39 31-05-2018

40 31-05-2018

41 31-05-2018

M 62,07 100

V 74,21 120

M 55,65 100

42 31-05-2018 V 73,91 100 JA

75,99 100

62,94 100 JA JA

100

JA JA

JA JA

JA

JA

JA

JA

JA

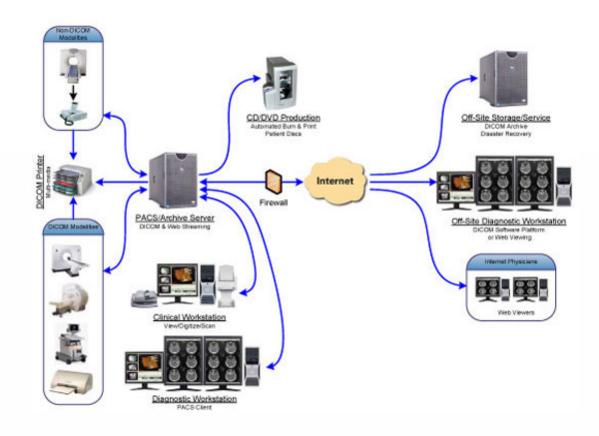
JA

JA

JA

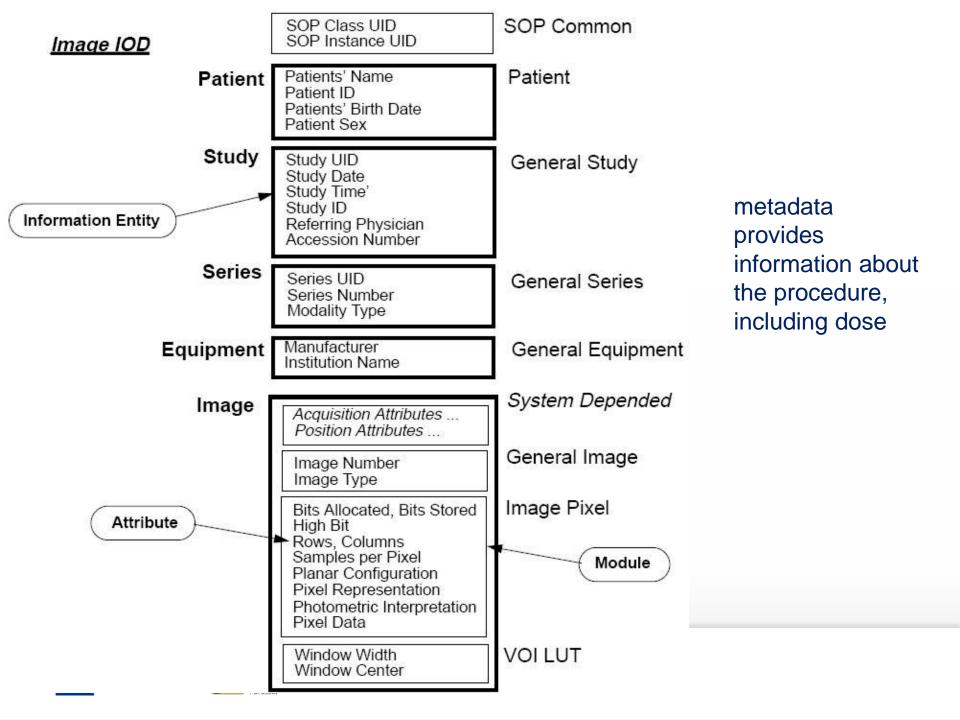
## REGISTRATION OF DOSES IN PATIENT IMAGES

Individual dose data are indicated on the console and archived in images (DICOM headers).







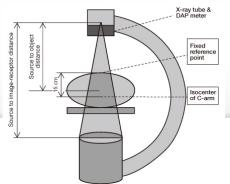


## DICOM HEADER

		· · · · · - · · · -	
0018 00	15 6	body_part_examined	CS   1   "CHEST"
0018 00	60 6	kvp	DS   1   "125.0"
0018 10	00 12	device_serial_number	LO   1   "966169249110"
0018 10	20 32	software_versions	LO   1-n   "3.1.2\PMS81.101.1.1 GXR GXRIM9.1"
0018 10	30 6	protocol_name	LO   1   "Thorax"
0018 10	50 6	spatial_resolution	DS   1   "0.143"
0018 11	10 6	distance_source_to_detector	DS   1   "1800.0"
0018 11	11 6	distance_source_to_patient	DS   1   "1746.0"
0018 11	50 2	exposure_time	IS   1   "3"
0018 11	52 2	exposure	IS   1   "1"
0018 11	53 4	exposure in uas	IS   1   "700"
0018 11	5e 4	image area dose product	DS   1   "O.23"
0018 11	64 12	imager_pixel_spacing	DS   Z   "U.143\U.143"
0018 11	66 2	grid	CS   1-n   "IN"
0018 14	00 64	acquisition_device_processing_descrip	otion   LO   1   "UNIQUE: S:200 L:4.0 FC d:1,8
0018 14	05 4	relative_x_ray_exposure	IS   1   "270"
0018 51	01 2	view_position	CS   1   "PA"
0018 70	50 16	filter_material	CS   1-n   "COPPER\ALUMINIUM"

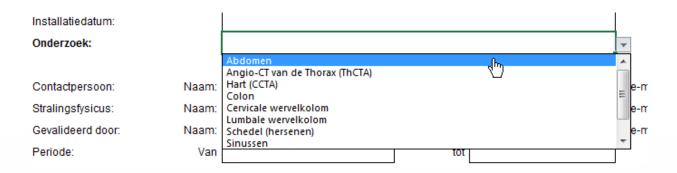








## COMPARE LOCAL DOSE TO DRL



VOLWASSENEN	c	TDI <sub>vol</sub> (mGy)	DLP (mGy.cm)						
VOLWASSENEN	Enkelv	oudig onderzoek	Enkelvo	oudig onderzoek	Volledig onderzoek				
Onderzoek	P25	DRN (P75)	P25	DRN (P75)	P25	DRN (P75)			
Abdomen	7	10	320	490	350	570			
Angio CT van de thorax	5	11	165	300	180	330			
Hart (CCTA)	10	25	140	290	190	480			
Colon	3*	4*	150*	200*	230	460			





## CAVEAT: DATA CLEANUP

## **Example: CT abdomen**

#### 34 different protocols used for 1 study description

- Wrong protocol selection
- Pediatric data
- Combined studies (thorax abdomen)
- Ultra-low dose protocol for follow up included
- → IMPACT on statistics
- → Data should be cleaned up before analysis



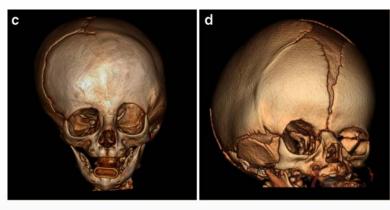


## WHAT IS THE INDICATION?

Today, technical exposure parameters (including dose) are selected based on indication

#### Example CT head

Diagnosis of craniosynostosis in children with cranial deformities



C. Ernst, Eur Radiol (2016)

Soft tissue information is not required

80 kVp; 8 mAs; CTDIvol = 0,9 mGy

Belgian DRL CT Head: CTDIvol = 45 mGy



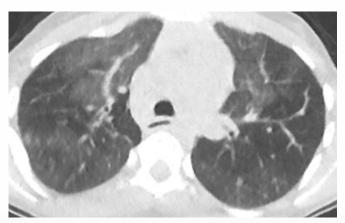


## WHAT IS THE INDICATION?

Today, technical exposure parameters (including dose) are selected based on indication

#### **Example CT chest**

Assessment of structural lung abnormalities in patients with cystic fibrosis (CF)



C. Ernst, Radiology (2014)

80 kVp; 4 mAs; CTDIvol = 0,09 mGy

Belgian DRL CT chest: CTDIvol = 3,5 mGy



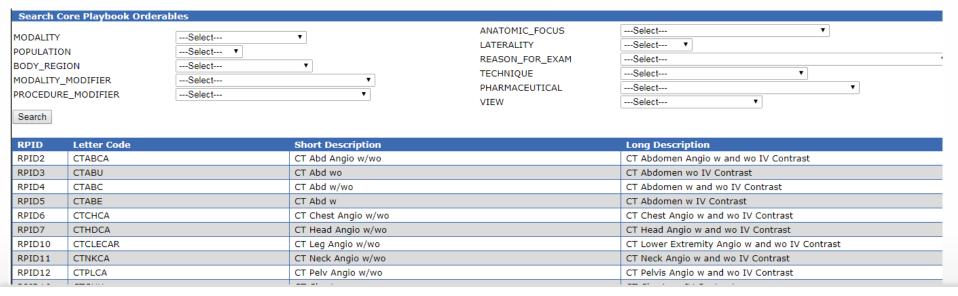


## STANDARD RADIOLOGY CODES

A standard system for naming radiology procedures, based on the elements which define an imaging exam such as modality and body part.

RPID code









## CAVEAT: DATA CLEANUP

## **Example: CT head**

	RadLex code	# patients	CTDIvol (mGy)
CT Hersenen (ALL)		1153	34,0
Hersenen axiaal	22	612	40,4
CT stroke (hersenen+perfusie+halsvaten)	96	458	25,1
Hersenen axiaal -/+ contrast	23	34	34,7
Flebo hersenen (sinus trombose)	7	49	21,1

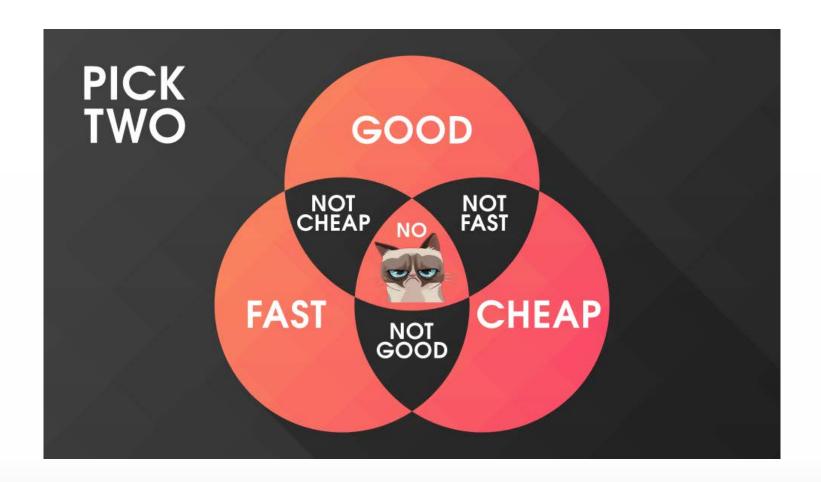


Belgian DRL CT Head  $CTDI_{vol} = 50 \text{ mGy}$ 





## PROJECT MANAGEMENT TRIANGLE







## DOSE MANAGEMENT

- Protocols with high dose compared to national and international reference values
  - Optimisation of higher-dose protocols
  - Uniformisation between systems
- New systems
- New techniques / changed protocols
- Check correct practices
  - CT positioning
  - Retakes

. . .





#### COMPARISON WITH NATIONAL AND INTERNATIONAL REFERENCE VALUES

Table 4.6. Typical patient dose levels (rounded) from gastroenterology and hepatobiliary procedures.

Procedure	Relative mean effective dose to patient	Relative mean radiation dose to patient*					Refere	nce'						
	0 mSv 35	to patient	Fluoroscopy time (min)	Entrance skin dose (mGy)	Dose-area product (Gycm <sup>2</sup> )	Effective dose (mSv)								
ERCP (diagnostic)		C,D	2-3	55-85	15	3-6	a,b							
ERCP (therapeutic)		E,F	5-10	179-347	66	20	a,b	÷						
Biopsy		C	n.a.	n.a.	6	1.6	a,c	TABLE 14: Summary of data on	other exam	inations a	nd interver	tional proced	dures (adults)	
Bile duct stenting		E	n.a.	499	43-54	11-14	a,c,d	Examination/procedure	Number			Mean of	Mean of	7
Percutaneous transhepatic		D	6-14	210-257	31	8.1	a		Hospitals	Rooms	Patients	room mean DAP	room mean fluoro. time	į
cholangiography			10.00										(seconds)	ì
Bile duct drainage		F,G	12-26	660	38-150	10-38	a,d,e					(Gy cm <sup>*</sup> )		
Transjugular intrahepatic		F,G	15-93	104-7160	14-1364	19-87	a,e,f	AICD	6	9	222	7	235	
portosystemic shunt creation								Angiography (Cerebral)	5	8	913	69	772	8
Transjugular hepatic biopsy		D	6.8	n.a.	34	5.5	f	Angiography (Mesenteric)	8	11	118	151	1009	7
ERCP, endoscopic retrograde	cholangio-pancreatograph	y; n.n., not availa	ble.					Angiography (Renal)	6	7	64	48	361	_7

10

10 18

12 19

Angioplasty (Femoral)

Angioplasty (Iliac)

Aortography (Arch) Arthrography (Hip) Dacryocystogram Electrophysiology

Embolisation (Uterine fibroid)

Embolisation (Varicocele)

ERCP (Diagnostic)

ERCP (Interventional)

Filter (Inferior Vena Cava)

Naso-gastric feeding tube

Pacemaker (Biventricular) Pacemaker (Temporary)

Percutaneous Transhepatic

Radio Frequency cardiac catheter

Radiologically Inserted Gastrostomy

Mean patient weight range 75-85 kg.

Retrograde pyelography

Stent (Biliary) Stent (Bowel)

Stent (Iliac artery) Stent (Superior Vena Cava)

Stent (Ureteric)

Thoracic spine Urodynamics

Right Heart Catheterisation

Cholangiography PTCA 2 stents\*

Percutaneous Endoscopic Gastrostomy

Oesophageal dilation

149 49

225 52

179

399

273 120

362

820

198 21

198

234 on

1761

815 52

2510

206

1238

803

23

13

19

588

1019

1715

625

154

270

233

664

138

653

1348

165

82

270

691

338

525

fluoro. time voltage (seconds)

74

72

71

71

70

69

70

70

74

81

71

75

(a) UNSCEAR, 2010; (b) Olgar et al., 2009; (c) Hart et al., 2002; (d) Dauer et al., 2009; (e) Miller et al., 2003a; (f) McParland, 1998.

#### Internationale referenties (HPA 2010)

ICRP 117 report, 2010





· DRN's bij volwassenen

				udig onderzoek		dig onderzoek
Onderzoek						
Abdomen	7	10	320	490	350	570
Angio CT van de thorax	.5	11	165	300	180	330
Hart (CCTA)	10	25	140	290	190	480
Colon	3*	4*	150*	200*	230	460
Cervicale wervelzuil	14	25	230	450	80	090
Lumbale wervelzuil	18	26	380	600	20	727
Schedel (hersenen)	35	50	640	900	- 80	686
Sinussen	2.5	6	35	80	22	350
Thorax	5	8	170	260	*	696
Thorax - abdomen	5	8.5	320	550	480	800





<sup>\*</sup> A, <1 mSv; B, 1-<2 mSv; C, 2-<5 mSv; D, 5-<10 mSv; E, 10-<20; F, 20-35 mSv; G, >35 mSv, based on effective dose.

### COMPARISON WITH NATIONAL AND INTERNATIONAL REFERENCE VALUES

		CT a			CT b			CT c		DRL CTE	)I enklel	DRL DL	P enkel	DRL DL	P totaal
	CTDI	DLP enkel	DLP totaal	CTDI	DLP enke	DLP totaal	CTDI	DLP enkel	DLP totaal	P25	P75	P25	P75	P25	P75
thorax	5.42	228.56	240.67	10.70	377.75	378.93	4.32	164.97		5.5	9	200	320		
abdomen	9.21977	519.78993	532.3206	11.88	614.40	634.45	7.31	395.77	425.65	7.5	12	350	600		
thorax-abdomen			753.34			750.35	5.05	278.77	420.23					550	960
CWZ	14.73939	397.18788	460.3429	28.37	569.09		10.61	231.63		16	30	280	490		
LWZ	26.96135	837.1366	838.3216	27.11	711.16	3	21.80	616.45	i e	20	30	400	650		
schedel				23.32	439.22	529.29	41.58	854.78	873.69	39	55	660	950		
sinussen	6.37	108.67		1.76	25.02	87.38				3	7	40	90		
hart CTCA	13.95857	246.8	255.3566				6.76	100.44	110.92	14	35	200	490		
angio CT thorax	8.3	402.30	528.3				2.22	89.75	156.96	6	15	180	410		
colon			501.9											230	530



Justification / optimization needed!



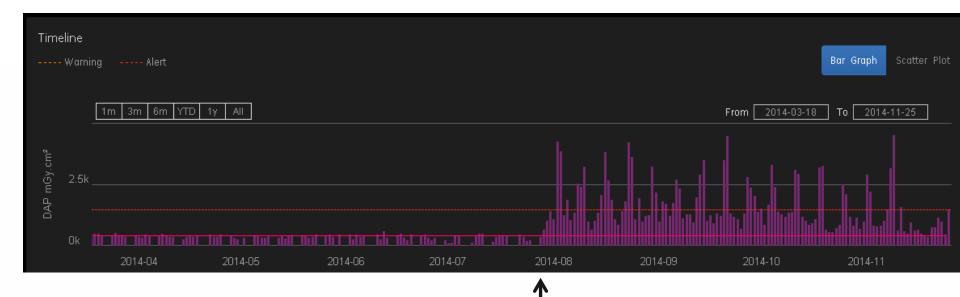




#### FOLLOW-UP ON NEW SYSTEMS

Daily dose UZ Brussel

Thorax PA+LAT



New RX system connected with dose registration



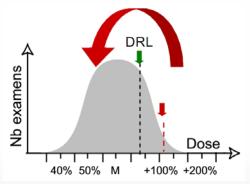


#### FOLLOW UP ON NEW SYSTEMS

	Room A	Room B	Room C	Room D	Room E	Room F
DAP (mGy.cm <sup>2</sup> )	506.37	352.16	468	297	58.05	2654
aantal procedures	7580	484	265	10	1482	1728

	national P75
national P25	(DRL)
530	1450





Conventionele radiologie bij volwassenen

On	derzoek	DRN in DAF	(cGy.cm²)
		25e p	75e p
Abdomen		120	330
Bekken face (Al	P)	170	450
Thorax PA		13	35
Thorax lateraal		40	110
Thorax aan bed		12	25
Lumbale	volledig (*)	7,5 <sup>(*)</sup>	21 <sup>(*)</sup>
wervelzuil	face	95	280
	profiel	200	500
Schedel	volledig	60	150
	face	25	60
	profiel	20	60





#### FOLLOW-UP ON NEW SYSTEMS

	Room A	Room B	Room C	Room D	Room E
DAP (mGy.cm <sup>2</sup> )	506.37	352.16	468	297	58.05
aantal procedures	7580	484	265	10	1482

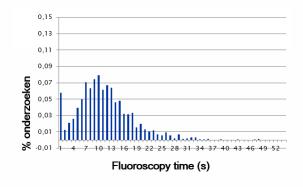
Room F	
2654	
1728	

	national P75
national P25	(DRL)
530	1450

Human

**Awareness** 

Reason?



**Equipment** 

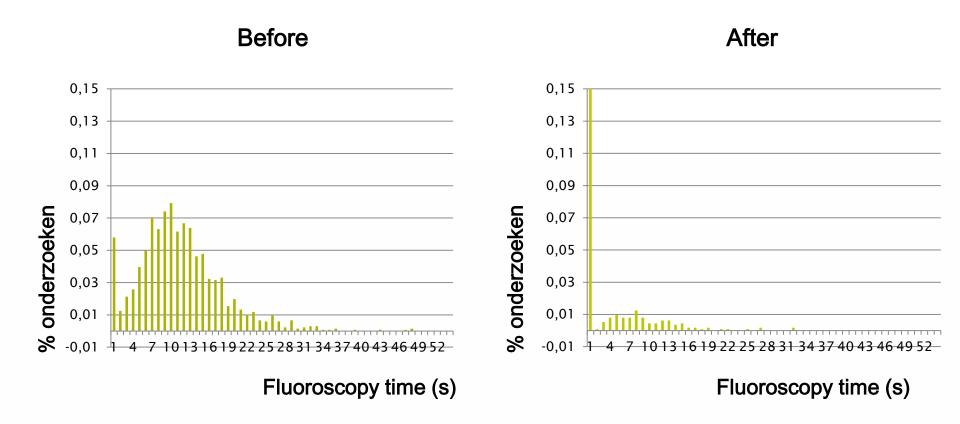
Fluoroscopy dose rate very high
Dose of 1 sec fluoroscopy = PA+lat radiography

→ Technical intervention





#### USE OF FLUOROSCOPY FOR POSITIONING



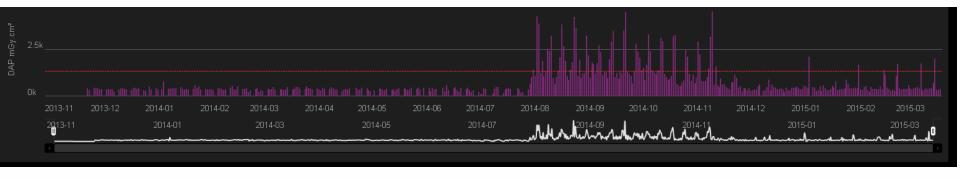
### **Awareness + Training**





#### 4 MONTHS LATER ...

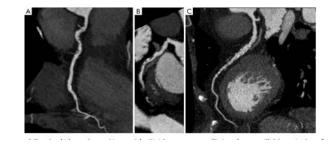
Mean dose over time in Radiology department for thorax PA + lateral



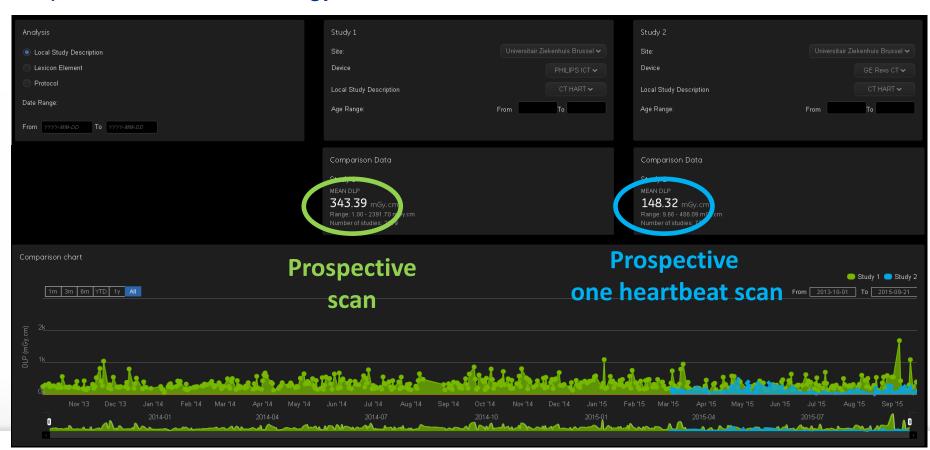




#### COMPARE CT SCANNERS



### Impact new CT technology for CCTA:







# CASE STUDY RX THORAX : #RETAKES ?

## 2 Month follow up

1377 thorax examinations

Standard of care: 2 images (PA and lateral)



- → individual comparison with PACS
- ▶ 192 DE opnames (separate protocol needed ?)
- ▶ 26 insp/exp opnames (separate protocol needed ?)
- ▶ 56 retakes (4%)







#### CONCLUSIONS

- Use the proper dosimetric quantity for your clinical practice
- Maintain dose values as low as reasonably achievable considering the clinical benefit of the procedure
- Work under radiation dose limits (for professionals) and use "Diagnostic Reference Levels" (DRLs) for patients. Avoid too high or too low doses.
- Avoid radiation injuries and establish a clinical followup for high dose procedures





#### TAKE HOME MESSAGE

A lot of technical tools are available to measure, record and evaluate patient doses from medical imaging



