





Protecting and improving the nation's health

DRLs and exposure monitoring in CT:

quantities, procedures, methods, international experience

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

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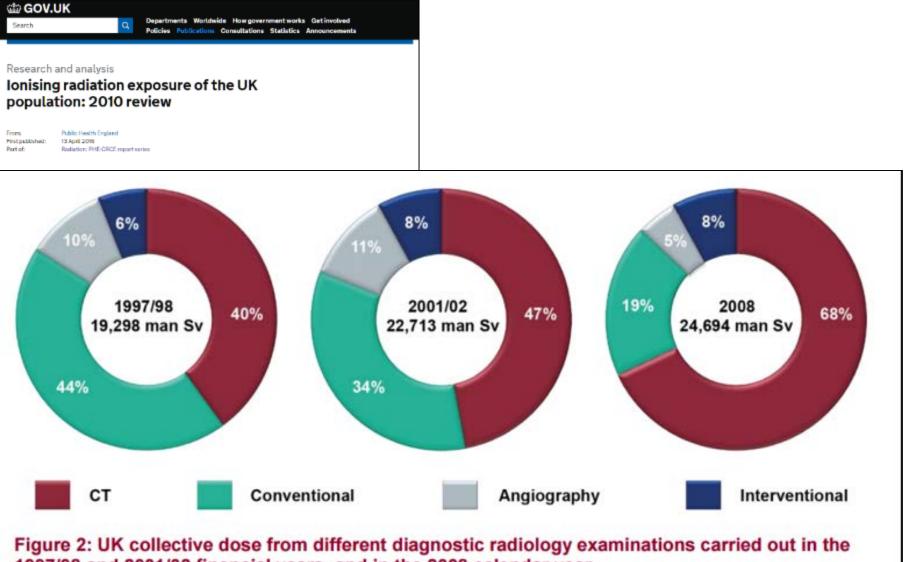
Outline

- CT Scanner
 - Fundamentals, Dose distribution
- Considerations for dose audit in CT
 - Dose index data (CTDI, DLP, SSDE)
 - How to get the data (manual dose monitoring systems)
 - Selection of exams
 - Selection of patients (size and numbers)
 - Relevant information to collect
 - Automatic Exposure control
 - Iterative reconstruction
 - Other things to consider SPR, contrast monitoring
- UK data



CT procedures deliver approximately 50% of the collective effective dose from medical and dental exposures in many countries, due to the relatively high-dose nature of CT procedures compared with other diagnostic imaging modalities (NCRP, 2009).

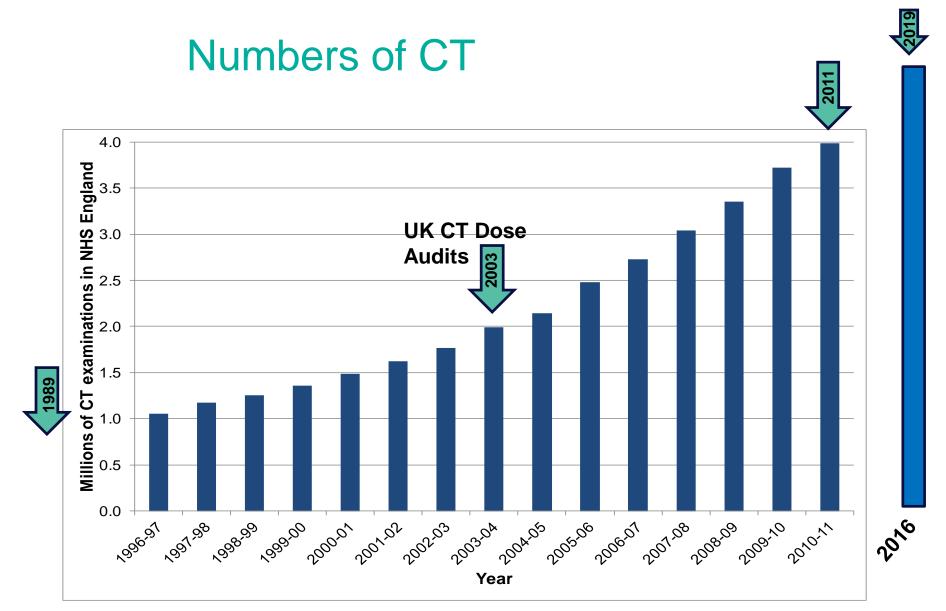
This contribution is increasing.



1997/98 and 2001/02 financial years, and in the 2008 calendar year

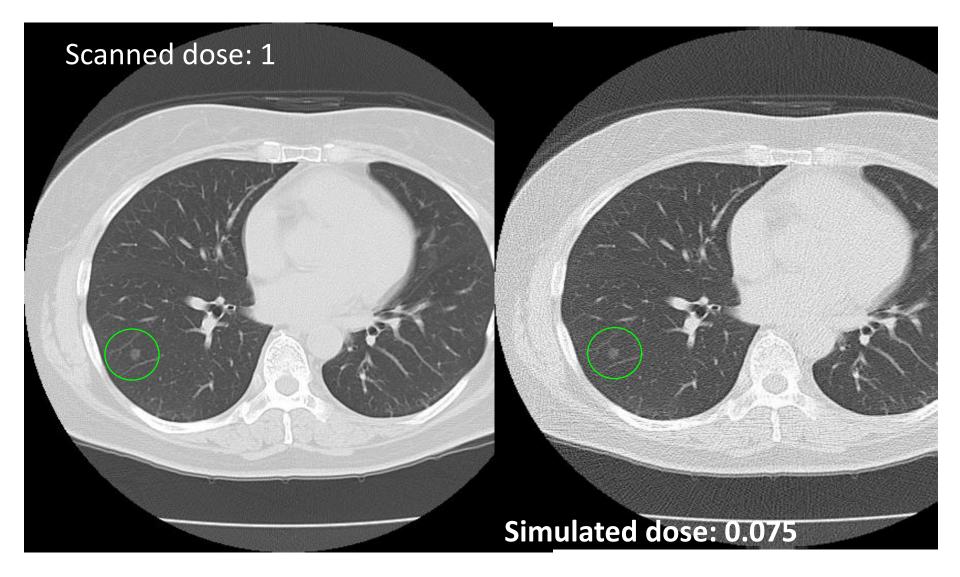
CT accounted for 68% of dose for radiology examinations in 2008

This is affected by level of dose and numbers of examinations



Annual numbers of CT examination performed in the NHS in England (Department of Health, 2011) (NHS England 2016)

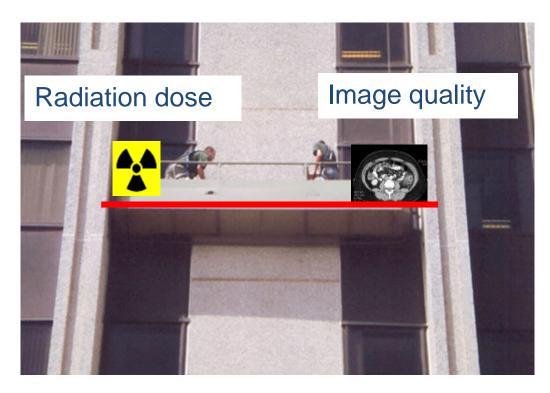
Image Quality in CT gets better and better with more dose



Images courtesy Y. Muramatsu, NCC Tokyo

Public Health CT Scanners – digital systems

- Detectors have high dynamic range –
- unbounded higher image quality for higher dose







Diagnostic Reference Levels

• All about ..

BENCHMARKING DOSES....

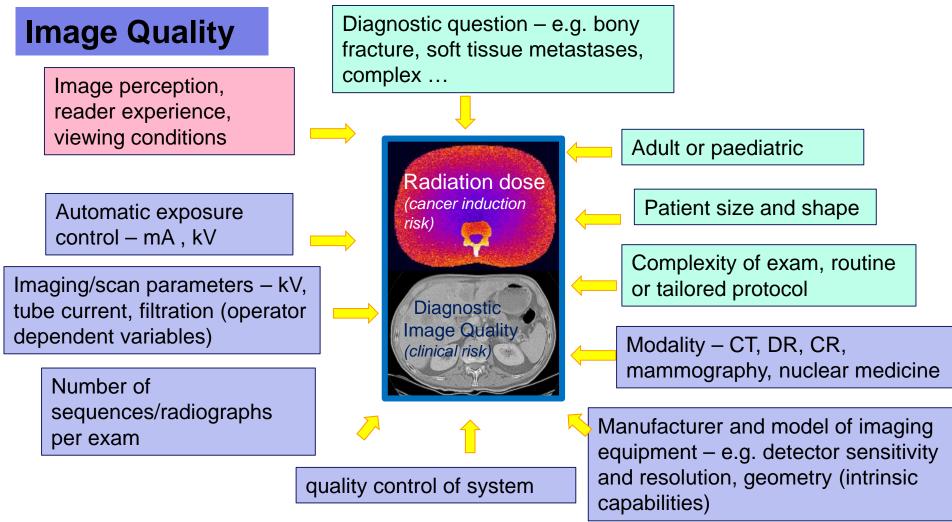
• From ... Dose Audits

• Using

STANDARD CONDITIONS AND BASIC STATISTICS

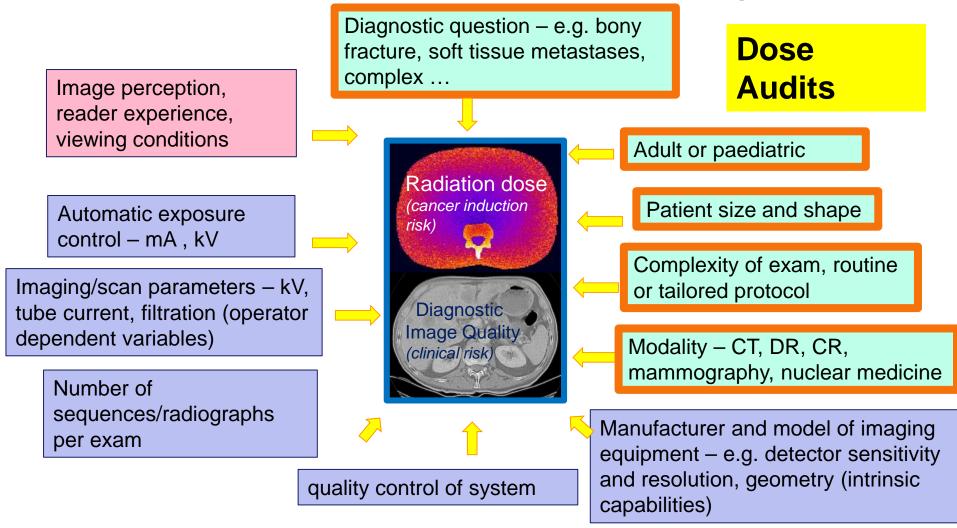
Factors influencing dose (and image quality)

A diagnostically acceptable image is the basic premise for DRLs



Factors influencing dose (and image quality)

For DRLs – some standardisation is required for a meaningful result ...





Dose Audits for DRLS

- Dose indicator (e.g. DAP,ESD or CTDI,DLP)
 - common examinations (e.g. chest CT) or high dose
 - Sample of standard size/weight patients
- Calculate the median^ value for each x-ray system, each exam

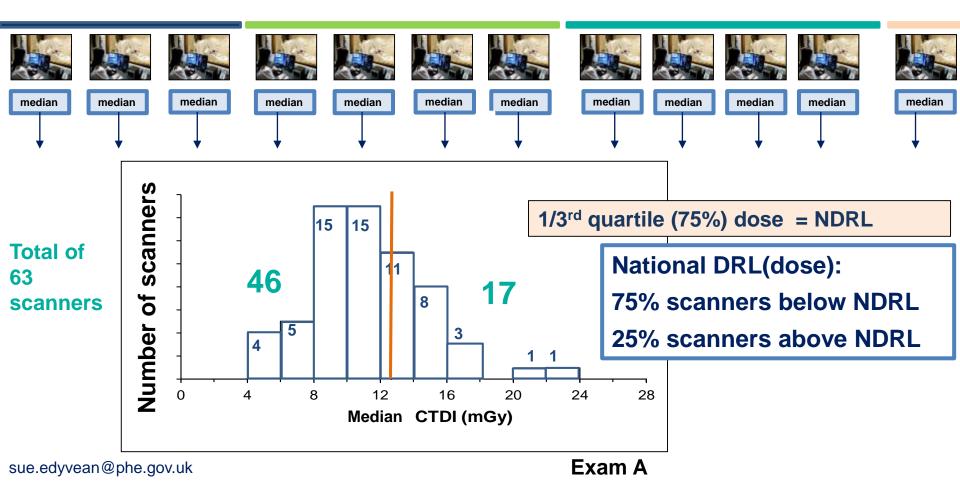


• ^UK previously used mean. UK currently ask for both: for retrospective comparison, and continue to future with median.

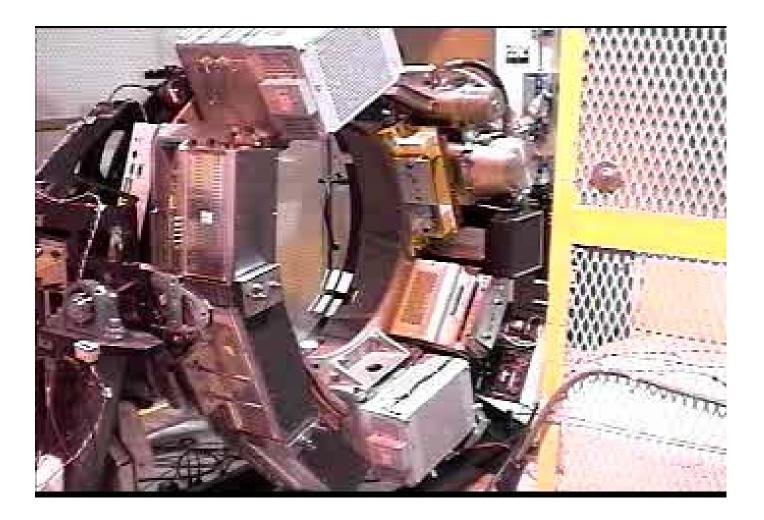


Two distributions of data

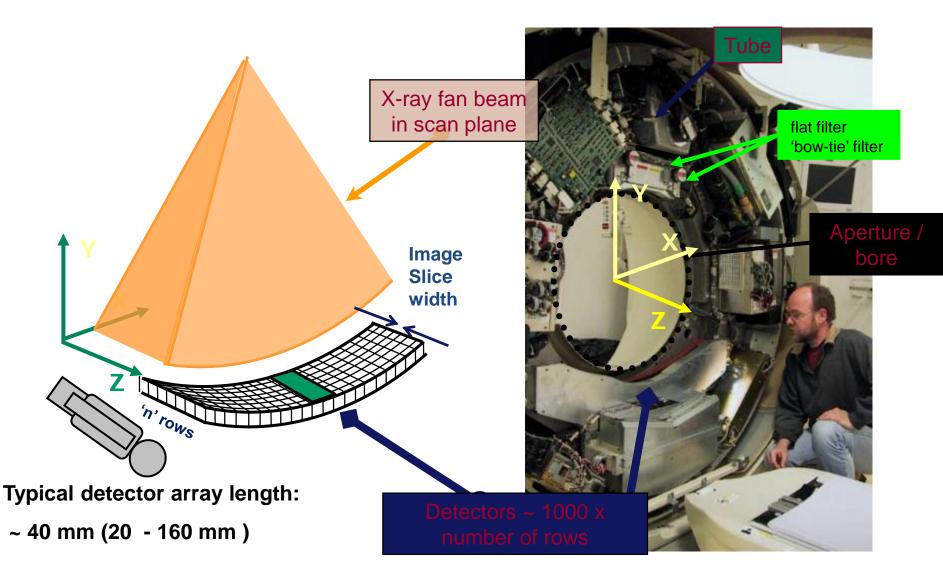
Distribution of Median Values from all scanners

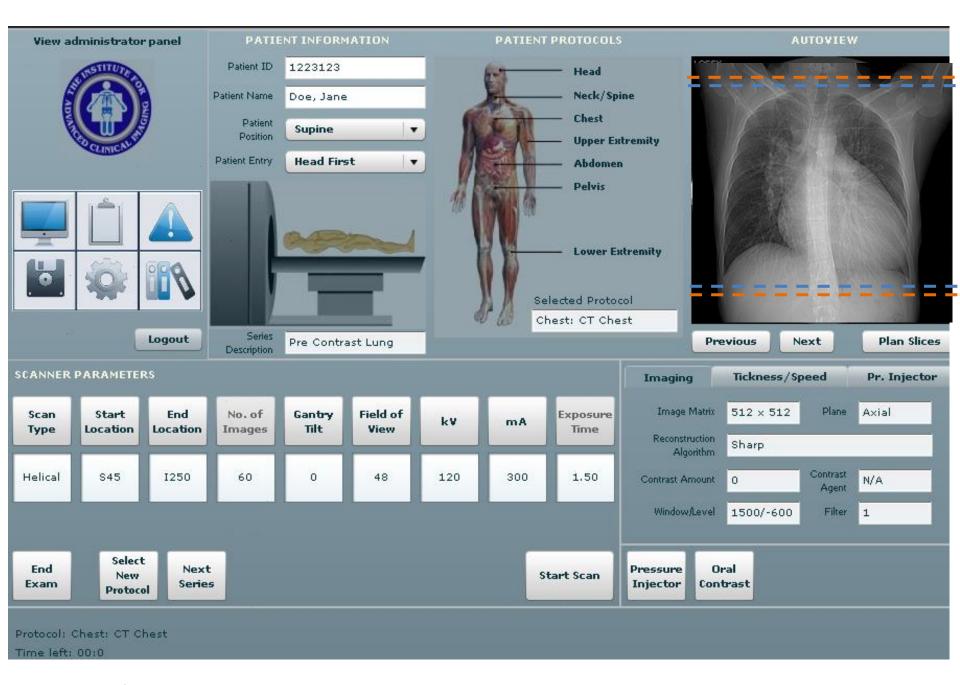


Technology



The Conventional MSCT Scanner



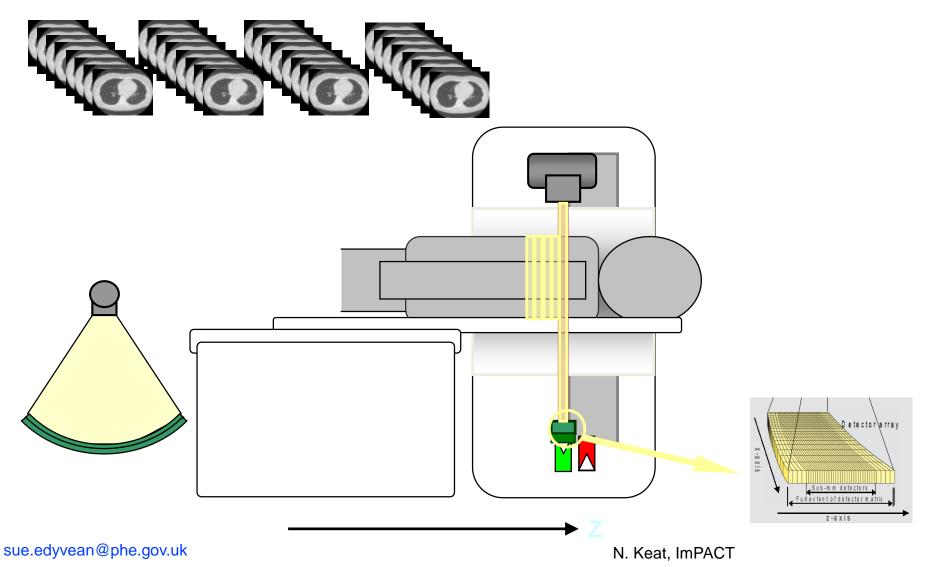


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http://www.iacionline.com/skins/userfiles/image/CTScanner_ScreenShot.png

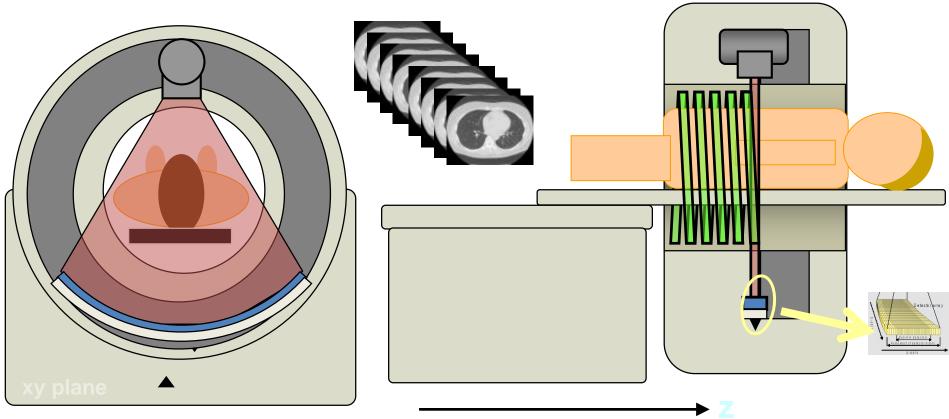
Axial scanning – 'step and shoot'

- Also known as sequential scanning



Helical (spiral) scanning

- Continuous gantry rotation + continuous table feed
- Multi-slice helical data used to form axial images



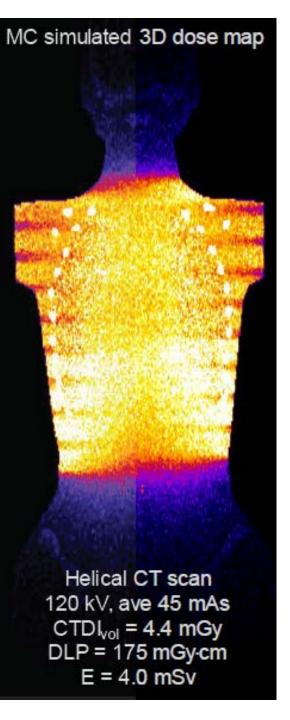
N. Keat, ImPACT

MSCT Examination -Dose Distribution in Z-Axis

Complex dose distribution

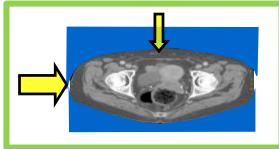
MC simulated dose map for a helical scan

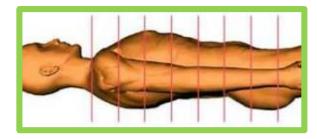
Courtesy Mika Kortesienmi



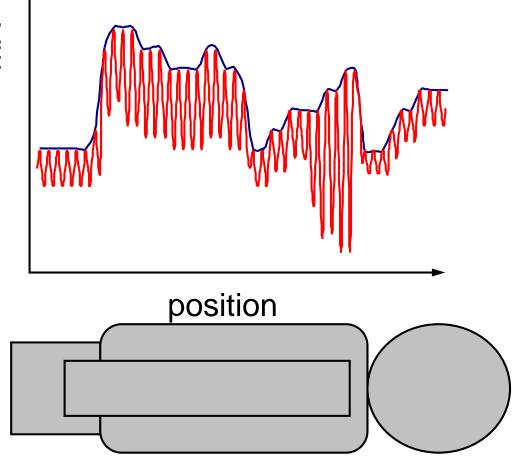
Automatic Exposure Control (AEC)



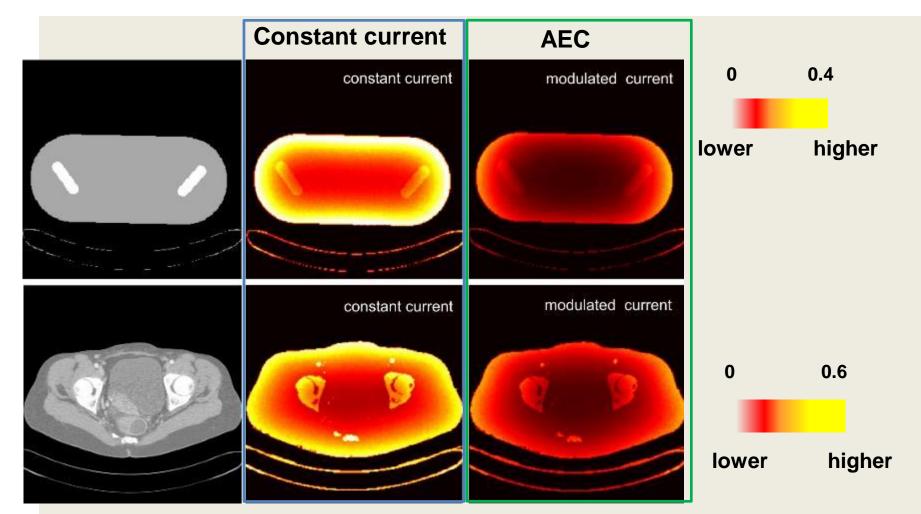


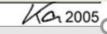






MSCT dose distribution in Scan Plane





Dose Metrics (Indicators) in MSCT

- CTDI mGy Computed Tomography Dose Index
- $MSAD_L$ mGy Multi-slice Average Dose (= $CTDI_L$)
- $D_0(L)$ mGy Cumulative dose = MSAD_L
- SSDE mGy Size Specific Dose Estimate
- DLP mGy.cm Dose Length Product
- ED mSv Effective Dose

$\label{eq:ctdlvol} \begin{array}{l} \text{CTDI}_{\text{vol}} \text{ and } \text{DLP used for setting } \text{DRLs} \\ \text{SSDE may be used to aide optimisation} \end{array}$

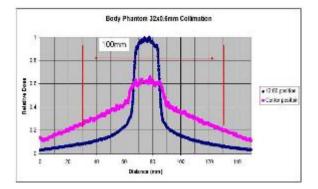


Quantities suitable for setting DRLs in CT

Quantity	Recommended symbols	Recommended unit	Closely similar qu	antity
Volume computed tomography dose index	CTDIvol	mGy	Volume CT air kerma index	(Cvol)*
Dose-length product	DLP	mGy.cm	Air kerma-length product	(Ркс)*

IAEA Web page

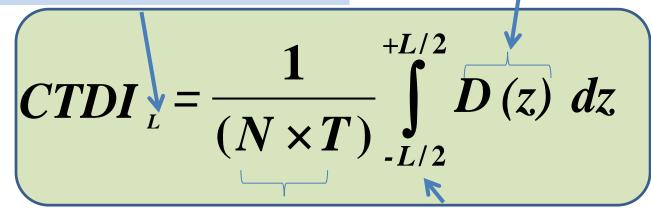
CTDI - general



A descriptor telling about the type of CTDI

(integration length, or medium measured in)







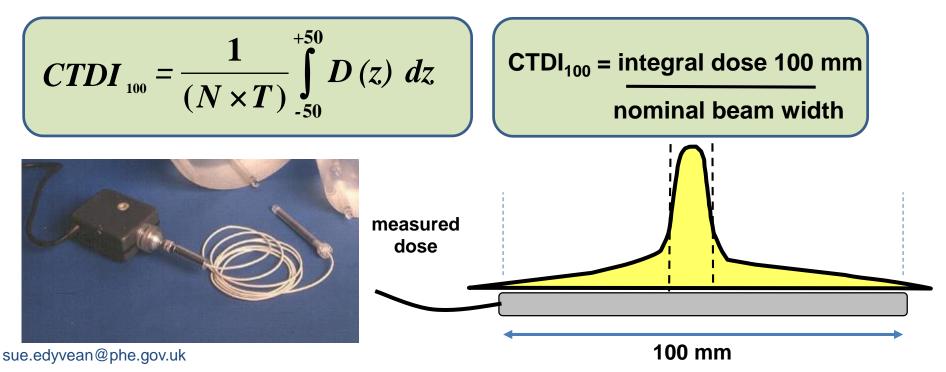
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The nominal beam width

Integral limits – how much dose we collect from the dose profile

CTDI₁₀₀

- 100 mm long ion chamber used
- Scan one rotation one 'dose slice'
- Dose from the profile is collected over 100 mm
- CTDI₁₀₀ is calculated: integral dose / nominal beam width



Weighted CTDI (CTDI_w)

- CTDI₁₀₀ measured in a Perspex phantom (quoted as dose to air)
 - 32 cm or 16 cm diam. (body, head)
 - Centre and periphery positions
- Cross-sectional average: CTDI_w = 1/3 CTDI_{100c} + 2/3 CTDI_{100p}



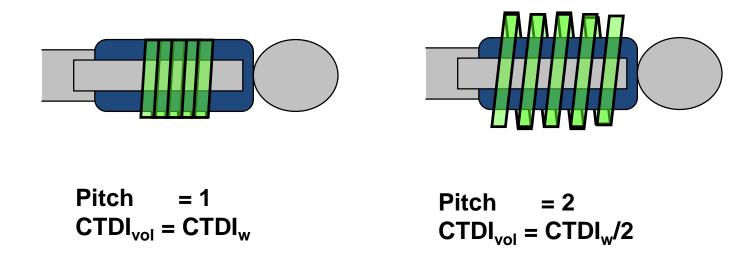
Manuf. data - tolerances 10-40%

Volume CTDI (CTDI_{vol})

- CTDI_{vol} takes account of exposure variation along z-axis
 - Accounting for pitch in the scan protocol

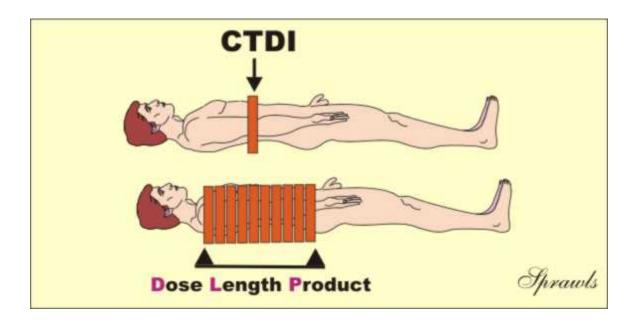
 $CTDI_{vol} = CTDI_{w} / pitch$

- CTDI_{vol} (axial scans) = CTDI_w x packing factor
- CTDI_{vol} ~represents average absorbed dose (x,y,z)



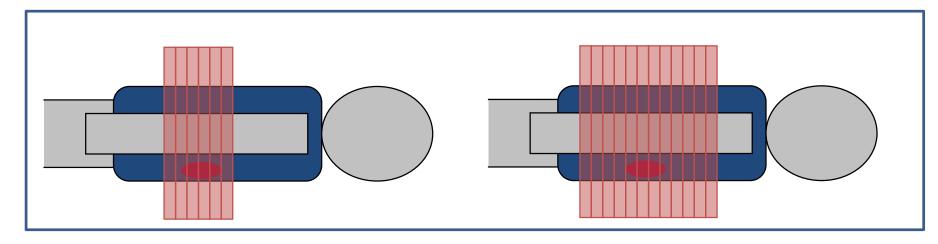
Dose length product (DLP)

- Dose descriptor used to indicate total absorbed dose
- Relates to risk
- DLP (mGy.cm) = CTDI_{vol} x scanned length (L).



Double imaged length – same mAs

 $CTDI_{vol} = same$ DLP = x 2



e.g. $CTDI_{vol} = 10 mGy$ DLP = 200 mGy.cm $CTDI_{vol} = 10 mGy$ DLP = 400 mGy.cm

CTDI and DLP – Dose Audits

- CTDI relates to cross-sectional scan parameters
 - Suitable for each sequence
 - DLP relates to clinical input wrt length of scan
 - Suitable for total exam, and each sequence if available

These may have

- same or similar CTDIvol,
- but will have different DLP

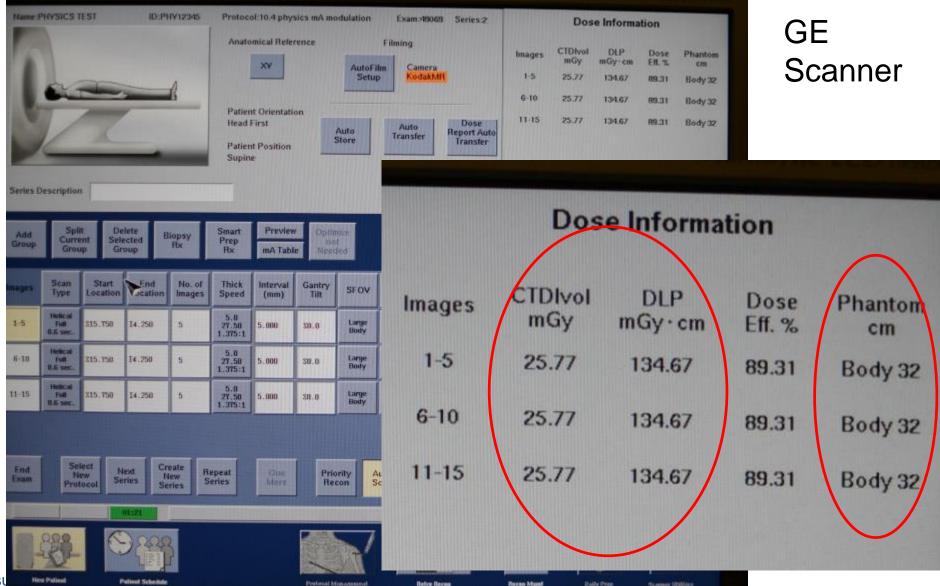




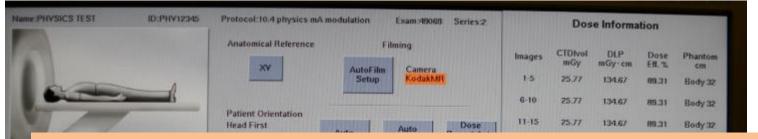
Each sequence:

 different average CTDIvol

CTDI_{vol} and DLP are indicated on the scanner



$\ensuremath{\mathsf{CTDI}_{\mathsf{vol}}}$ and DLP are indicated on the scanner

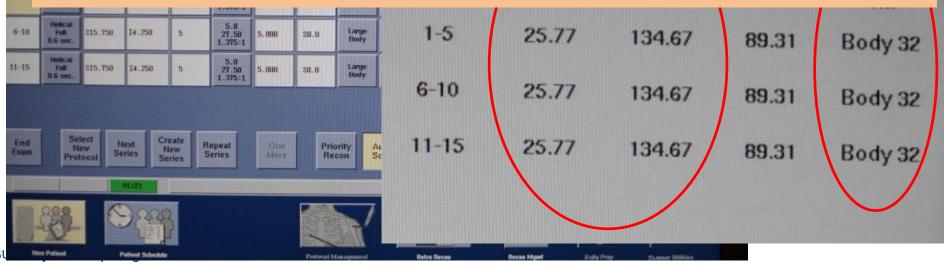


GE Scanner

Information will be given before and after the scan Where AEC is used the value presented will be the average value over the whole examination:

- Before the scan will be an estimate
- After the scan will be the actual

1-5



CTDI_{vol} and DLP are indicated on the scanner

Dose Report/Dose page – stored as an image

Birth Date : 1946.10.10	Age: 64Y				
Sex : M Weight(kg) : 96	Height(cm): 183.0				
Patient Comments :					
Study Date : 2011.08.18	Body Part : CORONARY ANG LOGRAM				
Requesting Department :					
Referring Physician : CT					
Reporting Physician : CF					
Operator Name : RD					
Total Image Number : 1257					
(Dose Information >>					
Total mAs : 7258	Total Scan time : 36.78				
CTDIvol(mGy) (Head) : -	(Body): 567.30				
DLP(mGycm) (Head) : -	(Body): 1735.90				
Contrast/Enhance Information >>					
Contrast Enhance ' CE					

Size Specific Dose Estimate (SSDE)

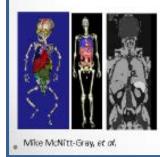
- Effective diameter (AAPM 204)
- Water Equivalent Diameter (D_w) (ААРМ 220)
- SSDE = $CTDI_{vol} \times f$

SSDE a dose index or estimate more representative for the patient size

Table 1D				
Dw	Conversion			
(cm)	Factor			
8	2.76			
9	2.66			
10	2.57			
11	2.47			
12	2.38			
13	2.30			
14	2.22			
15	2.14			
16	2.06			
17	1.98			
18	1.91			
19	1.84			
20	1.78			
21	1.71			
22	1.65			

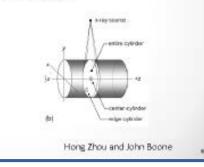


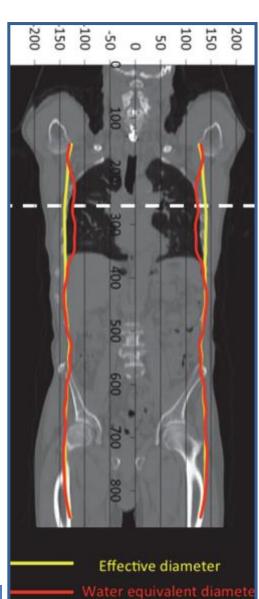






Cynthia McCollough et al.



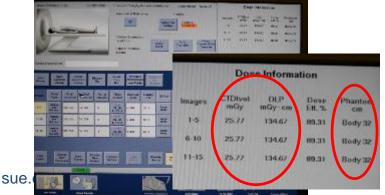


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AAPM Report 220

Dose Data - where is it?

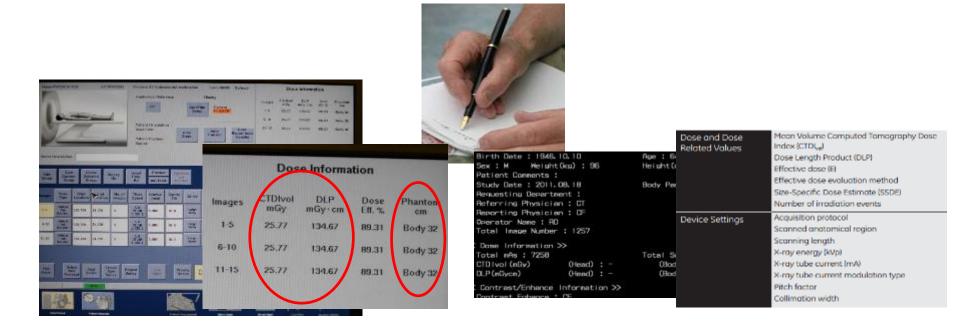
- Scanner
 - On the screen
 - Dose page (get as image or as digital data: optical character recognition (OCR))
 - In DICOM information: Radiation Dose Structured Report (RDSR)
- PACs (from dose page or DICOM RDSR)
- Dose Management System (from scanner or PACS, or RIS)
- RIS dose input manually from scanner (at the time of exam or after) (RIS – Radiology Information System)



Birth Date : 1946.10.10 Sex : M Melght(kg) : 96 Patient Comments : Study Date : 2011.08.18 Requesting Department : Referring Physician : CT	Ape : Br Height(c Body Per		Mean Volume Computed Tamagraphy Dase Index (CTDL _a) Dose Length Product (DLP) Effective dose (E) Effective dose evoluation method Size-Specific Dose Estimate (SSDE) Number of irradiation events
Reporting Physician : DF Operator Name : RD Total Image Number : 1257		Device Settings	Acquisition protocol Scanned anatomical region
:Dase Information >> Total mAs : ?258 CTDivol(nGv) (Head) : - DLP(nGyon) (Head) : -	Total Sc (Bod (Bod		Scanning length X-ray energy (kVp) X-ray tube current (mA) X-ray tube current modulation type
Contrast/Enhance information >> Contrast Echance : CE			Pitch factor Collimation width

Dose Data – How to get it ?

- Write / type into Excel
- Export electronically from: PACS, RIS, DMS
- Web based systems type info in



Importance of CTDI phantom Size

- For same mAs:
 - CTDI head phantom =~ twice CTDI body:
 - CTDI_{vol32cm} = 0.54 CTDI_{vol16cm}
- Important especially for
 - Paediatrics
 - cervical spine (neck scans)



(AAPM 2014)

Check phantom size used for CTDI value

American Association of Physicists in Medicine (AAPM). Use of water equivalent diameter for calculating patient size and size specific dose estimates (SSDE) in CT (task group 220). Maryland, USA: AAPM; 2014.

CTDI IEC standards, Phantom size

IEC 60601-2-44 Ed 3.1:

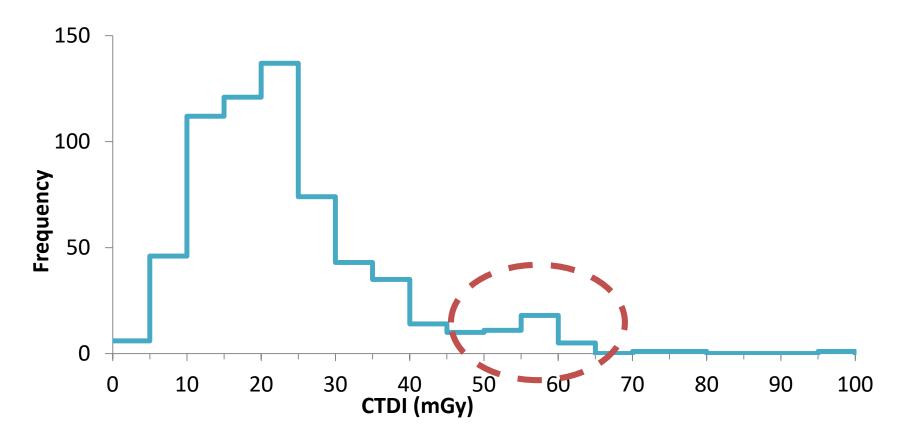
All bodies (adult and paediatric) 32 cm phantom

All heads (adult and paediatric) 16 cm phantom

Paediatric phantom specification given – Ed. 3.1 onwards SSDE to be introduced

CT Dose Disp	lay and Record	ing Require	ements in IEC 606	601-2-44		
IEC 60601-2-44 edition	date of standard	<u>clause</u>	Dose metrics to be displayed prior to scan	<u>Dose metrics to be</u> <u>displayed after the</u> <u>scan</u>	Dose Metrics to be recorded in <u>RDSR</u>	Accuracy of Dose display and <u>recording</u>
Ed. 2.0	June, 2001	29.1.103.3	CTDIw	n/a	n/a	n/a
Ed. 2.1	November, 2002	29.1.103.4	CTDIvol	n/a	n/a	n/a
Ed. 3.0	February, 2009	203.112	CTDIvol, DLP, phantom type (diameter)	CTDIvol, DLP, phantom type (diameter)	CTDIvol, DLP, phantom type (diameter)	n/a
Ed. 3.1	August, 2012	203.112	CTDIvol, DLP, phantom type (diameter)	CTDIvol, DLP, phantom type (diameter)	CTDIvol, DLP, phantom type (diameter)	The accuracy of the displayed and recorded values of CTDIvol and DLP shall be specified in the user manual.
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C-Spine (Fracture): Distribution of Scanner Median CTDI values (PHE CT 2011 Survey Report)



CTDI values for 7 scanners

CT Scanner		GE LIGHTSPEED VCT (64)				
Scanner ID	No of patients	Average CTDI	Median CTDI			
1	20	43.2	41.6			
2	20	17.2	16.4			
3	20	18.5	18.2			
4	20	16.4	15.7			
5	5	15.1	16.1			
6	20	32.4	30.6			
7	20	16.2	16.2			

IAEA_ICTP_Trieste_2019

J. Holroyd, PHE (data from PHE CT 2011 Survey)

Importance of CTDI phantom Size

- Cervical spine (neck)
 - <u>Head</u> and neck protocol 16 cm
 - Neck and <u>body</u> scan 32 cm
- Recent UK survey found both in use
 - and that most scanners use 32 cm. But the same model may be utilised differently (even in the same organisation)

Table 2. Choice of CTDI phantom used to calculate CTDI measurements by the scanners included in this survey

CTDI phantom	Number of scanners	
16 cm head	4	
32 cm body	69	
CTDI, CT dose index.	Holroyd JR, Edyvean S. Doses from cervical spine compution (CT) examinations in the UK. <i>Br J Radiol</i> 2018; 91 : 20170	



BJR

Table 9. Summary of cervical spine CT NDRLs in the UK: existing and proposed

IW astional DDL a	Quoted for 32	2 cm phantom	Quoted for 16 cm phantom		
UK national DRLs	CTDIvol (mGy)	DLP (mGy cm)	CTDIvol (mGy)	DLP (mGy cm)	
Existing	(15)	(324)	28	600	
Proposed	20	440	(37)	(815)	

AAPM, American Association of Physicists in Medicine; CTDI, CT dose index; DLP, dose-length product; DRL, diagnostic reference level; PHE, Public Health England.

Current value taken from the 2011 PHE CT survey is published for a 16 cm phantom. The actual proposed value from this study will be given for the 32 cm phantom. The conversion to the other phantom in each case is given in brackets, using the 0.54 factor from AAPM (2014).

Calibration / Verification of CTDI

- Manufacturers Specifications accuracy of actual CTDI – IEC +/- 20% or even 40%
- Values on the screen may be representative of that model, or made on the actual scanner at the factory.
- Only one collimation and set of scan parameters may have been be measured at subsequent tests
 - Other values obtained using specification correction factors for collimation, tube current, kV etc

Dose Information					
CTDIvol mGy	DLP mGy∙cm	Dose Eff. %	Phantom cm		
25.77	134.67	89.31	Body 32		
25.77	134.67	89.31	Body 32		
25.77	134.67	89.31	Body 32		
	CTDIvol mGy 25.77 25.77	CTDIvol mGy DLP mGy·cm 25.77 134.67 25.77 134.67	CTDIvol mGy DLP mGy·cm Dose Eff. % 25.77 134.67 89.31 25.77 134.67 89.31		

Calibration details – PHE Survey

Calibration Data (only if available)

Last measured CTDI _{vol} for this or a similar protocol (mGy):	
mAs used for the CTDI measurement above:	
Displayed CTDI _{vol} for the CTDI measurement above (mGy):	

Calibration / Verification of CTDI

PHE 2017 Cervical Spine (Neck) CT Survey

- In this survey, information was requested on the latest CTDIvol measurement made on the CT scanner.
 - Details were requested on the measured and reported CTDIvol values for the standard cervical spine protocol, or for the most similar protocol. See table 4 for results.
- As data were not corrected for error in the previous PHE CT dose survey, it was decided not to correct the data for this single exam survey.
 - However, analysis performed, without those scanners with a discrepancy greater than 10%, showed no significant effect on the final results, and therefore, this aspect did not need to be considered for the application of the final reference values.

• Table 4 summarises the information received.

- The vast majority of scanners had
 CTDI values measured within a few percent of the displayed values,
- with only four scanners having an error greater than ± 10%.

Number of scanners	27
Average error (%)	0
Standard deviation (%)	6
Minimum error (%)	-13
Maximum error (%)	17

Current UK National Patient Dose Audits Public Health England



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X-ray & Fluoro Pilot

UK Adult: plain X-ray; simple IR/fluoro surveys

Pilot		17 th	April 2019		
Mid-2018	2019	April	July	Sept	2020
Pilot		20 th Ma	arch 2019		
UK 4 th CT survey (adult) pilot UK 4 th CT survey (adult)					
			6 th Jur	ne 2019	
			-	IPEM/PHE	
	Computed Tomograph	ı y	UK 4 th CT	survey (pae) آ	diatric)

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Data collection survey

- By Excel spreadsheet, familiar design
 - PHE cervical spine CT audit
 - IPEM SPECT/PET CT and radiotherapy audits
- Distributed via
 - CTUG mailing list (ctug.org.uk)
 - UK Medical-Physics-Engineering mail list
 - SCoR (Society and College of Radiographers) website and newsletter



Next CTUG meeting: 3	rd October 2019
	Users Group will be held at The Studio in Birmingham traft programme, meeting details and booking form are meeting page.
UK Paediatric CT dose	survey
	sation working party, in collaboration with PHE, T dose survey in June 2019.
	he spreadsheet for dose data entry and guidance on can be found on the CTUC dose survey page
Fourth UK National CT	dose survey
	ounced on 23rd March 2019 their next review of doses o UK. This survey aims to collect protocol and patient T examinations.



Participants urgently needed for PHE's CT dose survey

22 July, 2019

More responses are urgently needed for Public Health England's (PHE) Fourth National CT Cose Survey

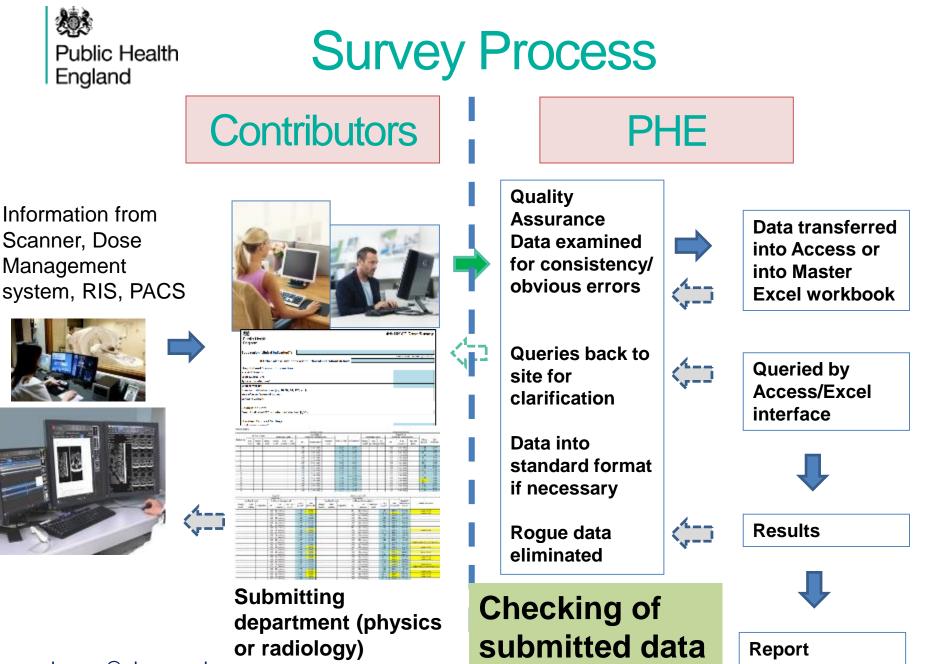
SCoR members are encouraged to subtrib data for any of the 13 examinations or other adult examinations that are rootinely performed and/or have the highest doses.

"It has been decided to extend the deadline date for submitting data until the end of September 2019 to give participants additional time. If you have date ready to submit, please do so as soon as you can which will help us with processing," says PHE

The data collection form and scanner help sheets are available on the CT users group website at Intp Dwww.ctug.org.sk/ctsurvey.html

For any queries and to submit date please email medicalisation/trass@plue.gov.uk







Patient Selection - Examination

- Selection of Exams for National Audit
 - High frequency (most common)
 - High dose
- Specify:
 - Anatomical region
 - Clinical reason for scan



PHE UK Dose Audits – Selection of Exams

NHS

Digital

Diagnostic Imaging Dataset

- Selection of Exams for National Audit
 - High frequency
 - High dose

Diagnostic Imaging dataset (NHS Digital/NHS England)

SNOMED-CT and/or NICIP RIS codes

SNOMED-CT Code Title	Count 2016	% of all
		exams
Computed tomography of entire head (procedure) (408754009)	1,209,740	26.1%
Computed tomography of thorax, abdomen and pelvis with contrast (procedure) (433761009)	539,640	11.6%
Computed tomography of abdomen and pelvis with contrast (procedure) (432370003)	434,450	9.4%
Computed tomography angiography of pulmonary artery (procedure) (419225001)	193,225	4.2%
Computed tomography of urinary tract (procedure) (419084009)	153,895	3.3%
Computed tomography of chest (procedure) (169069000)	151,370	3.3%
Computed tomography of thorax with contrast (procedure) (75385009)	144,820	3.1%
Computed tomography of thorax and abdomen with contrast (procedure) (429864007)	114,295	2.5%
Computerised tomography of chest with high resolution (procedure) (315941000000105)	101,755	2.2%

Preliminary survey



PHE Preliminary CT Dose Survey

Page 1
Dear Colleague,
Thank you for undertaking this preliminary survey to help inform the 4th review of doses from CT examinations in the UK.
The CT dose survey intends to look at the most frequent CT examinations, and/or those with the highest dose. The survey will only be considering adult examinations. A separate survey by IPEM in collaboration with PHE will be carried out to look at paediatric examinations.
The purpose of this preliminary survey is to help identify the examinations to request data for, as well as to get information on the level of detail that can be provided by different hospitals.
Please answer as many questions as possible. If you cover multiple hospitals, please complete a separate survey per hospital.
Thank you,
John Holroyd
Medical Dosimetry Group
Public Health England
medicalradiationdoses@phe.gov.uk

Preliminary survey results

Parameter	Automatic (%)	Manual (%)
Age	84	79
Patient diameter	15	33
Height	5	8
Weight	7	8
Size specific dose estimate (SSDE)	27	23
Water equivalent diameter, D _w	24	34

Could people supply us with this information? - and whether they had to obtain it manually (by weighing, or dimensions from images) or from a dose management/PACs/RIS system

2003 ()
Public Health
England

IAEA

CT Survey Workbooks: Colour

coding

Nublic Health	4th	UK CT Dose Survey
England		
Body region (clinical indication)*:		
If Other please give body	region (clinical indication) details:	Please select from drop down list
Hospital and Scanner Information		
Hospital Name*:		
Local system ID*:		
System manufacturer*:		
System model*:		
Number of detector rows (eg. 16, 32, 64, 128	3, etc):	
Year of manufacture of scanner:		
Software version:	Essential fields CT: blue	
Calibration Data		
Error of indicated CTDIvol when last checked	(+/- %)	
Standard Protocol Settings		
Local protocol name*:		

Pati	ent det	tails						This should distance for							This should l						
			t time of so					couch star							couch start						
			t time or st	an:	Scan	length (r	mm)	end.	tocol:			Scar	length (r	nm)	end.	rotocol:			50	an length (n	im)
Pati	ient No	Age (yrs)	Weight (kg)	Height (cm)	Imaged length	Start position	End position		Scan FOV (mm)	CTDI _{est} (mGy)*	DLP (mGy.cm)*	Imaged length		End position		Scan FOV (mm)	(mGy)*	DLP (mGy.cm)*	Imaged length	Start position	End position
	1																				
	2																				
	3																				
	4																				
	5																				
	6																				
	7																				
	8																				
	9																				
	10																				
	11																				
	12																				
ъ т	13																				



PHE UK Dose Audits – Selected Exams

Examination	Clinical indication	Suggested scan justifications that may use a similar exposure setup
Head	Acute stroke	head trauma, onset of headaches/facial pain, visual disturbances, aura/migraine, atypical seizure. Confusion, vomiting, slurred speech, limb weakness/worsening mobility. Existing aneurism. Previous surgery: CVA, evacuation of haematoma, biopsy
Paranasal sinuses	Sinus disease	Tumour, infection
Cervical spine (C-spine)	Fracture	head and neck injury. Fall/trauma/polytrauma. Previous vertebral tension. Neck pain or tenderness. RTC. Contact sports neck related injury
Neck, chest, abdomen and pelvis	Query Cancer	Query Lymphoma, lymphadenopathy, nodal disease
Chest	Ullery Lling cancer	Query cause of shadowing. Query lymphadenopathy. Previous lymph node enlargement. Bulky hilum (that persist on plain film). Abnormal CXR, pleural effusion
Chest – high resolution		Severe breathlessness, hypoxia, query parenchymal involvement. Subpleural ground- glass opacity
Chest and abdomen	Query Lung cancer	chest mass, abnormal CXR, shadowing, pleural effusion
Chest-abdomen-pelvis (CAP)	Query Cancer	Night sweats, weight loss, sepsis
CT pulmonary angiography (CTPA)	Pulmonary embolism	Pleuritic chest pain, decreased saturations, breathlessness. Sudden onset SOB. Previous surgery/PE
Abdomen and pelvis	Abscess	abdo pain, acute abdomen, weight loss, sepsis
Colonography/Virtual colonoscopy (VC)	Polyps/tumour	Anaemia, change of bowel habit, (do not include bowel cancer screening)
Kidney-ureters-bladder (KUB)	Stones/colic	Colicky pain, vomiting, previous calculus, haematuria
Urogram	Stones/colic or tumour	Query urological injury. Colicky pain, vomiting, previous calculus, haematuria. Query Urothelial tumour

Not included (but which were in 2011 survey): CT Angiography, Abdomen, Enteroclysis



Protocol names

StudyDescription	Short_Name	RPID
CT HEAD SURGICAL PLANNING WO CONTRAST	CT HEAD WO IVCON	RPID22
CT HEAD W/O	CT HEAD WO IVCON	RPID22
CT HEAD WO CONT	CT HEAD WO IVCON	RPID22
CT HEAD WO CONTRAST	CT HEAD WO IVCON	RPID22
CT Head Scan wo Contrast	CT HEAD WO IVCON	RPID22
CT Head w/o Con	CT HEAD WO IVCON	RPID22
CT Head w/o Con/Mag/Al	CT HEAD WO IVCON	RPID22
CT NEEDLE GUIDE BIOPSY NEURO	CT HEAD WO IVCON	RPID22
CT ORBIT EAR WO CONTRAST	CT HEAD WO IVCON	RPID22
CT ORBITS, SCREEN FOR MRI	CT HEAD WO IVCON	RPID22
CT Orb/Ear w/o	CT HEAD WO IVCON	RPID22
CT Orbit/Sinus MR Screen	CT HEAD WO IVCON	RPID22
CT Orbits or Ear wo Contrast	CT HEAD WO IVCON	RPID22
HEAD CT	CT HEAD WO IVCON	RPID22
HEAD WO	CT HEAD WO IVCON	RPID22
TCT HEAD	CT HEAD WO IVCON	RPID22
TCT HEAD W/O	CT HEAD WO IVCON	RPID22
TCT HEAD WO CONTRAST	CT HEAD WO IVCON	RPID22
TCT Head Scan wo Contrast	CT HEAD WO IVCON	RPID22
TCT Head wo Con	CT HEAD WO IVCON	RPID22
TCT Head wo Con/Mag/Al	CT HEAD WO IVCON	RPID22
TCT Orbits or Ear wo Contrast	CT HEAD WO IVCON	RPID22

List of the 19 exam names used at one institution for noncontrast head CT

Public Health England

Clinical Reason for Scan

Data is requested for the examinations listed below See the 'scan regions'

PHE CT protocol	Clinical indication
Head	Acute stroke
Paranasal sinuses	Paranasal sinuses
Cervical spine (C-spine)	Fracture
Neck, chest, abdomen and pelvis	Query Cancer
Chest	Query Lung cancer
Chest – high resolution	Interstitial lung disease
Chest and abdomen	Query Lung cancer
Introduction Guidelines P	Protocol guidance Scan Regi

Head	
Paranasal sin	uses
Cervical spine	e (C-spine)
Neck, chest, a	abdomen and pelvis
Chest	
Chest – high i	resolution
Chest and ab	
Chest-abdom	en-pelvis (CAP)
CT pulmonary	y angiography (CTPA)
Abdomen an	d pelvis
Colonograph	y/Virtual colonoscopy (VC)
Kidney-urete	rs-bladder (KUB)
Urogram	

PHE UK CT Protocols for National CT Dose Audit (Adults)

PHE CT protocol	Clinical indication
Head	Acute stroke

Clinical Reason for Scan

 Other similar reasons
 Suggested scan justifications that may use a similar exposure setup

 head trauma, onset of headaches/facial pain, visual disturbances, aura/migraine, atypical seizure. Confusion, vomiting, slurred speech, limb weakness/worsening mobility. Existing aneurism. Previous surgery: CVA, evacuation of haematoma, biopsy

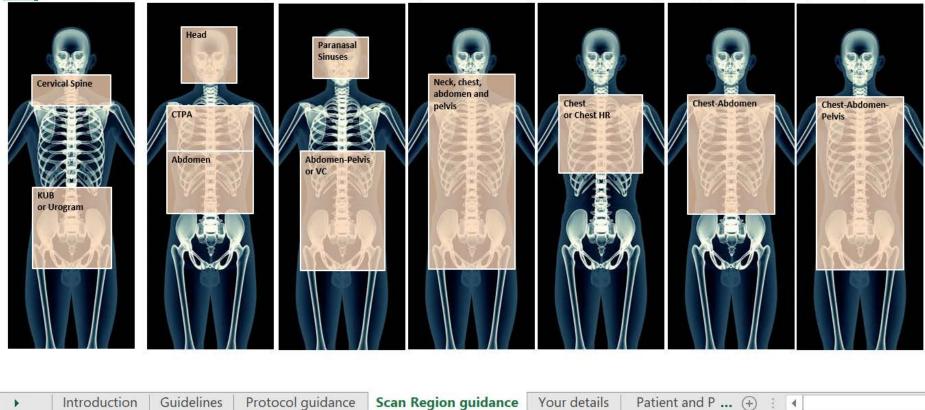
Key words for RIS or dose management search

Keywords for electronic searches (eg. on a RIS or dose management	
system)	
Stroke, CVA, haemorrhage	

Public Amatomical Scan Region Guidance

Rublic Health England

4th UK CT Dose Survey



The images below give an approximation of the start and end positions for the 13 examinations requested for this dose survey. These should be used as a guide only - please provide data for your clinical practice.

IAEA_ICTP_Trieste_2019



Guidance Notes

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Public Health	۱
England	

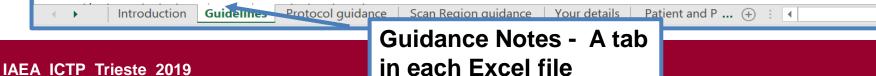
122

4th UK CT Dose Survey



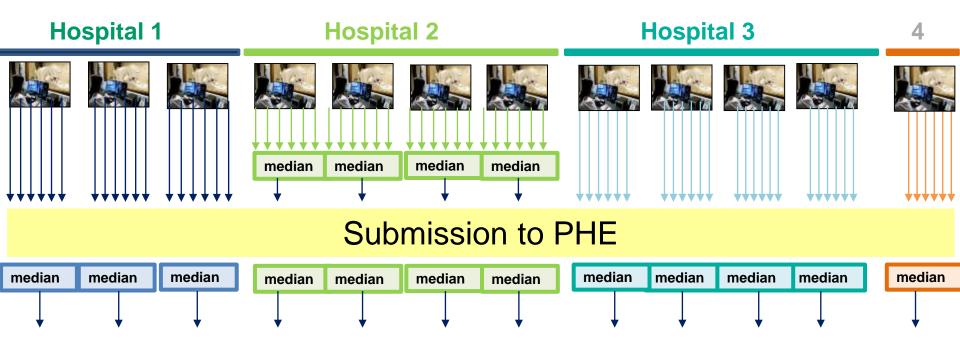
Guidelines

- 1. The data collection form may be completed by scientific, clinical or administrative staff. It is recommended that the scientific integrity of the data is checked by a medical physics expert (MPE) prior to submission.
- 2. Data may be acquired prospectively or retrospectively. For retrospective data, please ensure the data are still representative of current scanning techniques and protocols. Ideally data will be chosen from the previous year, but no older than 2017. Data from local dose surveys may be used if appropriate.
- 3. Only data from clinically acceptable scans should be included (ie. QA scans or scans where repeat exposures were necessary should be excluded).
- 4. Dose data is likely to be found within a dose management system, the images on PACS/other DICOM store, on the dose record page where available, or on the scanner console after the scan. The 'Protocol guidance' page provides details of typical examinations to search for.
- 5. There are also 'help sheets' available for different scanners which show you where to find the various parameters on your scanner.
- 6. Please supply patient weight information wherever possible.
- 7. No patient identifiable data should be included in your submission.
- 8. For each scanner and examination please supply data for as many patients as possible with a minimum of 20 different patients, but ideally at least 100 patients. There is no upper limit.
- 9. Patients should be selected who are considered a 'standard' size, ie. exclude patients who are atypically small or large. As a guide a weight range of 50 90 kg can be used.
- 10. Only data for adult patients should be submitted to this survey. For the purposes of this survey, an adult is anyone 16 years or older. For paediatric data, please refer to the IPEM/PHE paediatric CT survey forms.
- 11. Data is being collected for 13 different examinations. Please also submit data for other exams that you commonly perform



UK National DRLS

- Hospitals send either
 - Individual patient data or
 - Summary mean and median^ data from own audit



^asked for mean (for retrospective comparison), and median (ICRP recommended approach) for this and future surveys

\$\$\$ PHE 4th UK Survey – patient data Public Health England

82.5	Essential fie	lds			
戀			4th UK CT Dose Survey	Acquisition 1 details	
Public Health	(blue)			CTDI phantom size (cm) (i.e. 16	cm head or 32 cm b
England	(101010)			Is Automatic Exposure Control	
-				AEC name (e.g. AutomA, ZDOM	
Body region (dinical indication	1*:			AEC setting type (e.g. ref noise	
- ,			Please select from drop down list	AEC setting value (e.g. SD 7.5, r	
If Other please give	e body region (clinical indicatio	n) details:		minimum mA for AEC (where a	
				maximum mA for AEC (where a	
Hospital and Scanner Informati	ion			mA where AEC is not used:	
Hospital Name*:				is iterative reconstruction used	2*
Local system ID*:				Iterative recon type (e.g. ASIR,	-
System manufacturer*:				Iterative recon value (e.g. ASIR	
System model*:				Rediation beam collimation	- Collimated beam
Number of detector rows (eg. 16, 32	, 64, 128, etc):			Addated again commuter	- Number of slices:
Year of manufacture of scanner:					- Detector size (mr
Software version:				Is Automatic tube voltage selec	tion used? (eg. Carel
				If no, Fixed Tube voltage (kV):	
Calibration Data				Tube rotation time (s):	
Error of Indicated CTDIvol when last	chacked (al. 56)			Primary image slice thickness (r	nm):
End of indicated croniol when ast	enecked (47-36)			Scan field of view (SFOV) (mm):	
				Reconstruction field of view (Di	FOV) (mm):
Standard Protocol Settings				Axial or helical?	
Local protocol name*:				Pitch (where applicable):	
Number of scan acquisitions* (e.g. 1	contrast & 1 non-contrast scan = 2 ac	quisitions):		Primary Reconstruction algorith	um or kernel (e.g. B3
				is contrast used?	

Acquisition 1 details									
CTDI phantom size (cm) (i.e. 16 c									
Is Automatic Exposure Control (A	AEC) used?*								
AEC name (e.g. AutomA, ZDOM,	CARE Dose 4D, SureExpose)								
AEC setting type (e.g. ref noise in	ndex, reference mAa, etc):								
AEC setting value (e.g. SD 7.5, re	f mAs 200]:								
minimum mA for AEC (where app	plicable):								
maximum mA for AEC (where ap	plicable):								
mA where AEC is not used:									
is iterative reconstruction used?	•								
Iterative recon type (e.g. ASIR, SAFIRE, iDose, AIDR):									
Iterative recon value (e.g. ASIR 40%, SAFIRE 3, IDose level 4):									
Radiation beam collimation	- Collimated beam width (mm):								
	- Number of slices:								
	- Detector size (mm) (e.g. 0.625,0.6):								
Is Automatic tube voltage selecti	ion used? (eg. CarekV)								
If no, Fixed Tube voltage (kV):									
Tube rotation time (s):									
Primary image slice thickness (m	m):								
Scan field of view (SFOV) (mm):									
Reconstruction field of view (DFG	DV) (mm):								
Axial or helical?									
Pitch (where applicable):									
Primary Reconstruction algorithm	n or kernel (e.g. B30; FC17; Std)								
Is contrast used?									

Patient det	tails						/This should	i ka ika						This should be th							
Patient No	Age (yrs)	t time of sc Weight (kg)	an: Height (cm)	Imaged	i langth (n Start position	End	distance from the star and.	rum the 4. to couch	tocol: Scan POV (mm)	стрі _{на} (mGy)*	DLP (mGy.cm)*	Imaged	Start position	distance from the centh start to co end.	e Hith	srotocol: Scan FOV (mm)	CTDI _{ve} [m@y]*	DLP (mSy.cm)*	can length (n Start position	nm) End position	Submit
1 2 3 4 5																					by patient
6 7 8 9 10																					(no ID info)
10 11 12 13 14																					



Summary dose data from local audit

No of Patients	Mean Age at time of scan (yrs)	Mean Body Mass (kg)	Mean Total DLP* (whole scan)	Median Total DLP* (whole scan)	Comments on the data collection method (eg. inclusion criteria, data analysis method)

					Acquisition 1				
Mean CTDI _{vol} (mGy)*	Standard deviation	Median CTDI _{vol} (mGy)*	25th Percentile	75th Percentile	Mean DLP (mGy.cm)*	Standard deviation	Median DLP (mGy.cm)*	25th Percentile	75th Percentile

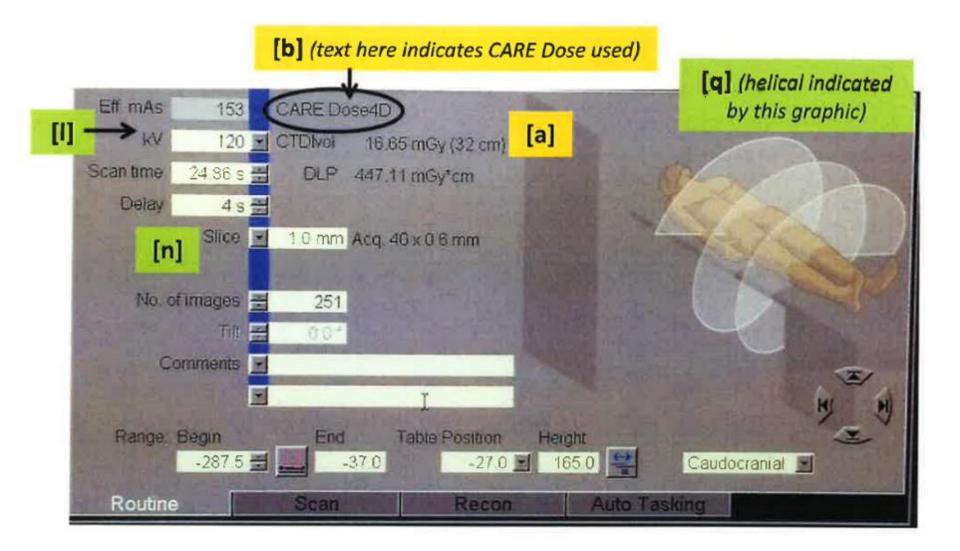
Acquisition 2											
Mean CTDI _{vol} (mGy)*	Standard deviation	Median CTDI _{vol} (mGy)*	25th Percentile	75th Percentile	Mean DLP (mGy.cm)*	Standard deviation	Median DLP (mGy.cm)*	25th Percentile	75th Percentile		

Or by summary data from local audit – for each system

Protocol details: scan details

Acquisition 1 details	See notes on scanner specific help shee	
CTDI phantom size (cm) (i.e. 16 cm hea	[a]	
Is Automatic Exposure Control (AEC) u	[b]	
AEC name (e.g. AutomA, ZDOM, CARE	[c]	
AEC setting type (e.g. ref noise index,	[d]	
AEC setting value (e.g. SD 7.5, ref mAs	200):	[e]
minimum mA for AEC (where applicat	le):	[f1]
maximum mA for AEC (where applical	ble):	[f1]
mA where AEC is not used:	[f2]	
Is iterative reconstruction used?		
Iterative recon type (e.g. ASIR, SAFIRE	[g]	
Iterative recon value (e.g. ASIR 40%, S	[h]	
Radiation beam collimation	- Collimated Beam width (mm):	[i]
	- Number of slices:	
	- Detector size (mm) (e.g. 0.625,0.6):	[k]
Is Automatic tube voltage selection us	sed? (eg. CarekV)	
If no, Fixed Tube voltage (kV):		[1]
Tube rotation time (s):		[m]
Primary image slice thickness (mm):		[n]
Scan field of view (SFOV) (mm):		[0]
Reconstruction field of view (DFOV) (r	nm):	[q]
Axial or helical?		[q]
Pitch (where applicable):	[r]	
Reconstruction algorithm or kernel (e	[s]	
Is contrast used?		
Anatomical landmarks for start and	Start point (e.g. base of skull)	
end points	End point (e.g. vertex)	

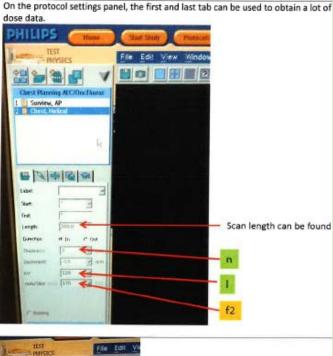
Siemens scanner - help sheet

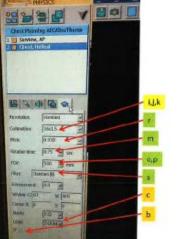


Philips CT scanners – help sheet

These instructions are based on an older Brilliance Big Bore scanner. New software versions may vary from guidelines.

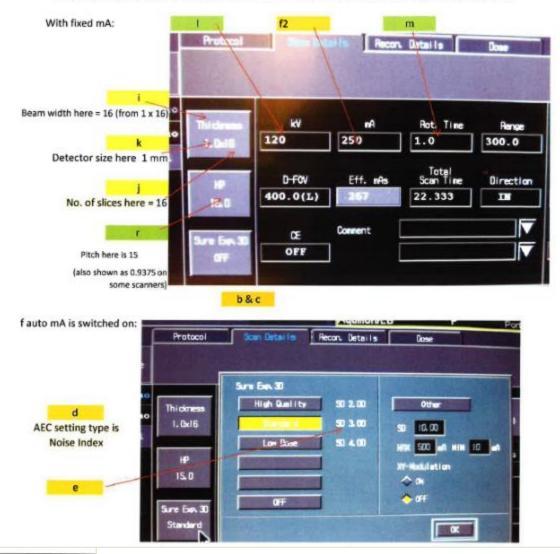
Prospective data collection (during patient scan):



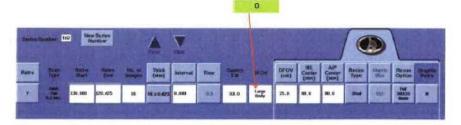


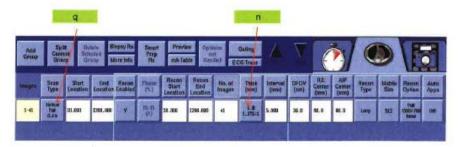
Canon / Toshiba scanners - help sheet

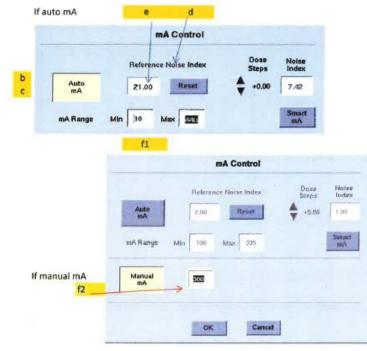
This is based on an old Toshiba LB. Newer scanners may look different but terms should be the same or similar.

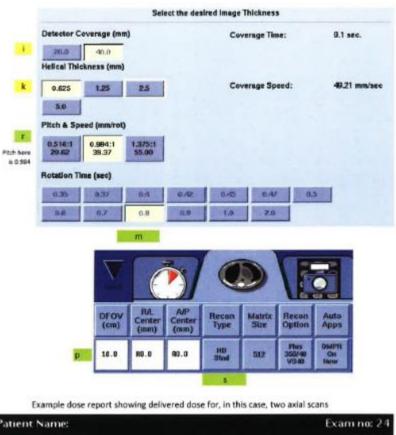


GE scanners - help sheet









Patient	Name:			Examino: 24			
Accessio	an Numb	er:	Nov 08 2011				
Patient	D: Dose	SR	Discovery CT750 HD				
Exam D	escriptio	in:					
		Dose R	eport				
Series	Түре	Scan Range (mm)	CTDivol (mGy)	DLP (mGy-cm)	Phantom cm		
1	Axial	\$0.000-\$57,500	93.37	560.24	Head 16		
1	Axial	\$60.000-\$135.000	63.88	511.06	Head 16		
		Total	Exam DLP:	1071.30			

a

Total mAs if available & switched on will be shown on the dose report. (Not shown here) If not available leave this part of the data sheet blank.

Patient Selection – size and numbers of patients

- Size of patient in sample
- Numbers of patients in sample

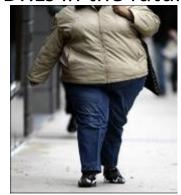




Dose Audits - Patient size

- Usually/previously specify data collected from
 - 70 kg +/- 20 kg (ie 50 90kg)
- So that the mean value of the weight is
 - 70 kg +/- 10 kg (or even +/- 5 kg)
- Of course standard weight is not 70 kg
- Weight often not available
- Now ICRP not so specific 'standard'
 - Causes problems for inter-comparison of DRLs
 - Good reason to have weight based DRLs in the future

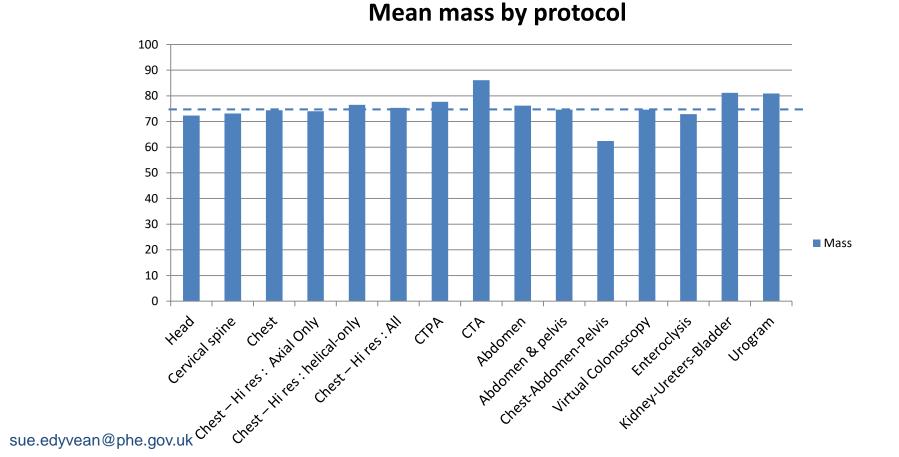






PHE 2011 Survey Data - Individual Patients

- Mean mass = 75 kg
- Max: CTA = 86 kg, Min: CAP = 62 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

A general accepted approach with large scale data sets is to remove the top 5% and bottom 5% of doses values

Large Scale Data – all weights

- E.g. from Radiology Information system (RIS) (with manual dose index data input), or PACs or Dose management systems
- Outliers can be removed easily e.g. removing top and bottom 5% of data

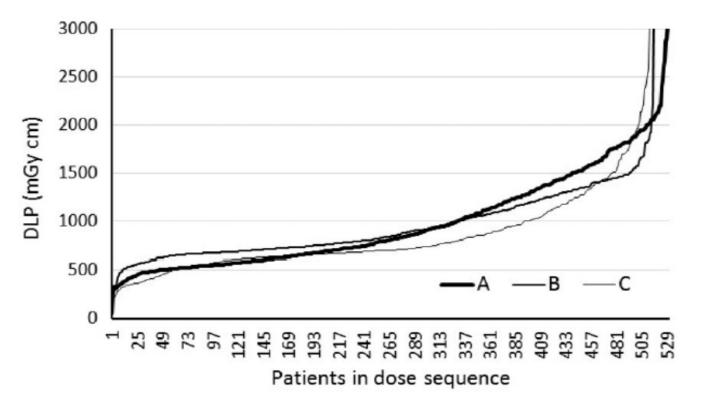


Fig. 2.1. Examples of data on dose-length product (DLP) for chest-abdomen-pelvis scans on three computed tomography (CT) scanners operating under automatic tube current modulation plotted sequentially in terms of increasing DLP (Martin, 2016). Outliers can be identified readily and omitted from the data analysis. Martin 2016, and ICRP 135⁷²

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VALIDATION OF A LARGE-SCALE AUDIT TECHNIQUE FOR CT DOSE OPTIMISATION

T. J. Wood^{1,*}, A. W. Davis¹, C. S. Moore^{1,2}, A. W. Beavis^{1,2,3} and J. R. Saunderson^{1,4} ¹Radiation Physics Department, Hull and East Yorkshire Hospitals NHS Trust, Queen's Centre for

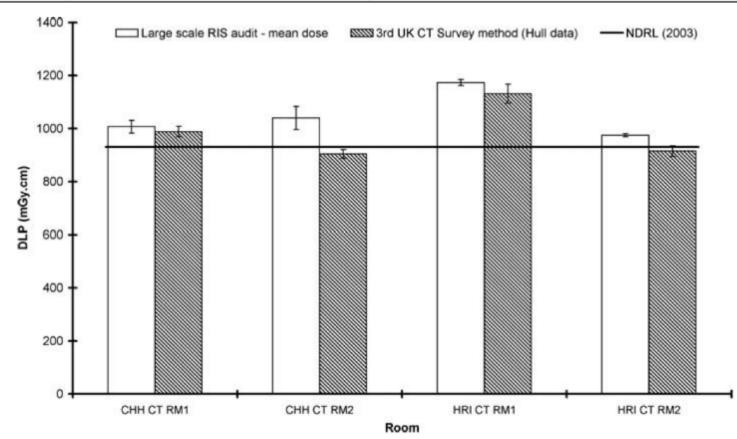
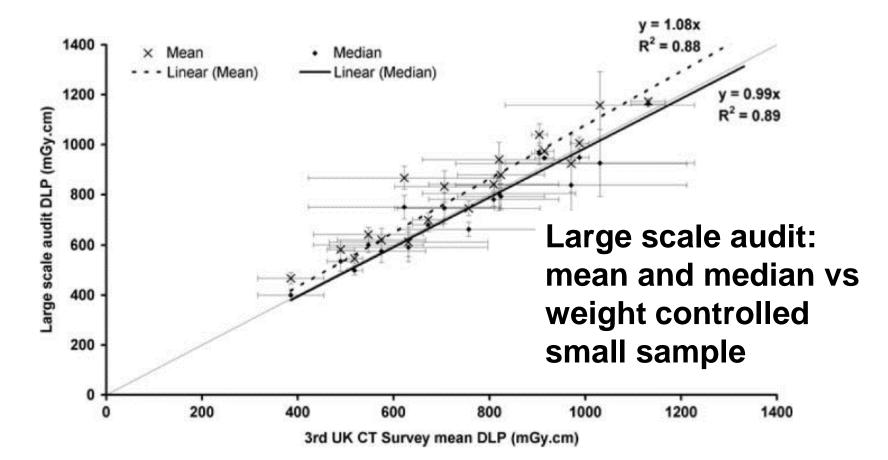


Figure 1. A comparison of the mean CT head DLPs determined using Dosalyzer with the third UK CT survey data, for each of the four radiology CT scanners in the Hull and East Yorkshire Hospitals NHS Trust. Error bars are defined by two times the SEM⁽⁸⁾, and the national DRL is indicated by the solid line (derived from the 2003 review of CT doses⁽¹³⁾).

VALIDATION OF A LARGE-SCALE AUDIT TECHNIQUE FOR CT DOSE OPTIMISATION

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ESTABLISHING LOCAL AND REGIONAL DRLs BY MEANS OF ELECTRONIC RADIOGRAPHICAL X-RAY EXAMINATION RECORDS

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The objective of the paper is to demonstrate that patient dose audits may be undertaken at the local and regional levels by employing electronic examination records contained in Radiology Information Systems (RISs) that have been collected, analysed and managed by modern IT systems. The resulting mean and third quartile values obtained may then be used to establish local and regional dose reference levels (DRLs) as part of an optimisation strategy. The method involved the collection of roughly 1.3 million radiographical examination records stored in hospital RIS over a 3-y period from 10 hospital sites in the north of England. These were analysed according to the process employed in the national patient dose (NPD) audits undertaken every 5 y in the UK. Data processing and analysis methods are described that are suitable for handling very large data sets quickly and efficiently. Because RIS data involve manual data entry it may be susceptible to data entry errors. Therefore, a comparison of results obtained from both RIS and DICOM generated data was first of all undertaken in order to 'calibrate' the RIS-based method and demonstrate its accuracy. The results obtained from this comparison indicate that the RIS-based examination records provide patient dose distributions with an equivalent statistical accuracy compared with those employing DICOM data and, therefore, may be employed in patient dose audits in order to establish both local and regional DRLs for use in patient dose management and optimisation strategies.

RIS (Radiology Information system)

Example of summary Data

- Total exam and DLP only

PHE CT Protocol	RIS Name		No of Patients	Mean Age at time of scan (yrs)	Mean Total DLP* (whole scan)	Median Total DLP* (whole scan)	Standard deviation
Abdomen and pelvis (Abscess)	Abdomen and pelvis (Abscess)	СТ	34	51.38	510.3	489	222.29
Cervical spine (C-spine) (Fracture)	Cervical spine (C-spine) (Fracture)	СТ	341	58.10	166.3	153	69.48
Chest (Lung cancer)	Chest (Lung cancer)	СТ	69	57.13	247.6	222.8	106.93
Chest-abdomen-pelvis (CAP) (Cancer)	Chest-abdomen-pelvis (CAP) (Cancer)	СТ	133	57.71	566.0	509	224.60
1	,	1 '		,	,		
CT pulmonary angiography (CTPA) (Pulmonary embolism)	CT pulmonary angiography (CTPA) (Pulmonary embolism)	СТ	54	57.17	267.2	264	69.12
Head (acute stroke)	Head (acute stroke)	СТ	2246	55.60	830.7	818.4	132.85
Other	CT Brain Volume (allegro)	СТ	717	50.08	900.3	947	213.62
Other	CT Spine Lumbar	СТ	464	54.77	211.7	184.5	107.70
Other	Angio Intracranal/Venogram Cerebral	СТ	573	52.38	728.2	744.2	88.45

Usually only get DLP data and total exam information from RIS

In the UK – since IRMER 2000 (UK law) radiographers input dose index data into RIS system



PHE 2019 survey - CT submissions to date



	This survey	2011 survey	
Number of Hospitals	60	127	A200/ of
Number of Scanners	115	182^	^30% of installed
Number of local audit spreadsheets	677	189	base
Number of patient spreadsheets	421	682	
Number of patients	413,257	46,938	

COMPUTED TOMOGRAPHY

CT dose survey in adults: what sample size for what precision?

Stephen Taylor¹ • Alain Van Muylem² • Nigel Howarth³ • Pierre Alain Gevenois⁴ • Denis Tack⁵

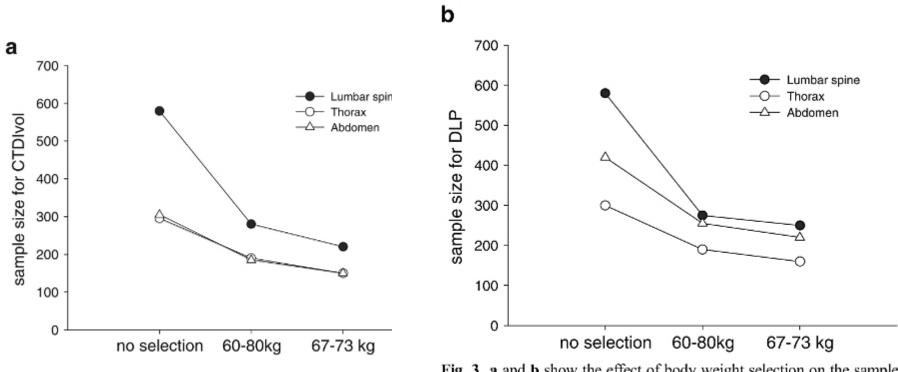
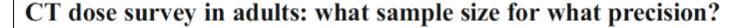


Fig. 3 a and b show the effect of body weight selection on the sample size required in center A to achieve CI95/med < 10 % for the thorax, abdomen and lumbar spine when using CTDIvol and DLP as the data sources, respectively

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sue.edyvean@phe.gov.uk

COMPUTED TOMOGRAPHY



Stephen Taylor¹ • Alain Van Muylem² • Nigel Howarth³ • Pierre Alain Gevenois⁴ • Denis Tack⁵

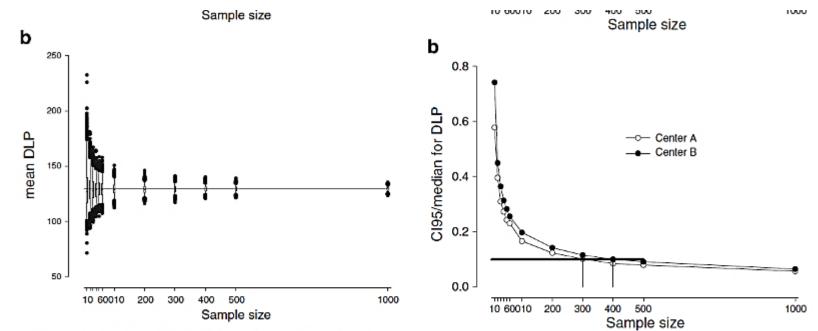


Fig. 1 a-b. show the sampled distribution of mean CTDIvol and DLP, respectively, for acquisitions in the thorax (2000 samples) as a function of sample size. For each sample size, the *box* represents the inter-quartile range and the *whiskers* represent the 95 % confidence interval; the *closed circles* are the values above percentile 97.5 and under percentile 2.5. Inside each *box*, the *horizontal line* is the median. The *dashed horizontal line* is the CTDIvol or DLP mean of the whole population of

Fig. 2 Thorax - a and b show the 95 % confidence interval for center A (*open circles*) and center B (*closed circles*) in percentage of the median as a function of the sample size, using CTDIvol and DLP as the data sources, respectively. Vertical lines corresponds to the sample sizes ensuring CI95/med < 10 %

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CT

PHE 4th UK CT Survey

LOS: 4th UK CT Dose Survey Public Health England Guidelines 1. The data collection form may be completed by scientific, clinical or administrative staff. It is recommended that the scientific integrity of the data is checked by a medical physics expert (MPE) prior to submission. 2. Data may be acquired proceedingly <u>ea ancura tha data ara ctill ran</u>r Please supply patient weight information wherever possible. current scanning te 5. from local dose surv No patient identifiable data should be included in your 6. Only data from clini submission. should be excluded 4. Dose data is likely to For each scanner and examination please supply data for as 7. page where availab many patients as possible with a minimum of 20 different examinations to sea patients, but ideally at least 100 patients. There is no upper 5. There are also 'help scanner. limit. Please supply paties 8. Patients should be selected who are considered a 'standard' No patient identifia 7. size, ie. exclude patients who are atypically small or large. As For each scanner ar but ideally at least 1 a guide a weight range of 50 - 90 kg can be used. Patients should be: guide a weight range of 50 - 90 kg can be used.

10. Only data for adult patients should be submitted to this survey. For the purposes of this survey, an adult is anyone 16 years or older. For paediatric data, please refer to the IPEM/PHE paediatric CT survey forms.

Dose Audits – Numbers of data and patient size

- Small data sample (manual methods of data collection):
 - 20 30 samples
 - Record and standardise patient size
- Large sample (automatic systems of data collection):
 - median size generally prevails

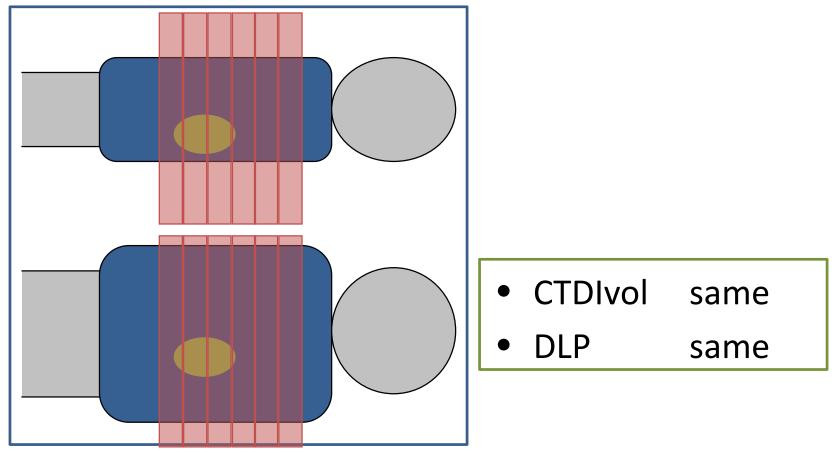




High 'dose' (CTDI) value may just mean you have scanned large patient, It does not necessarily mean *high dose* to the patient

Larger Patient Size – same CTDI

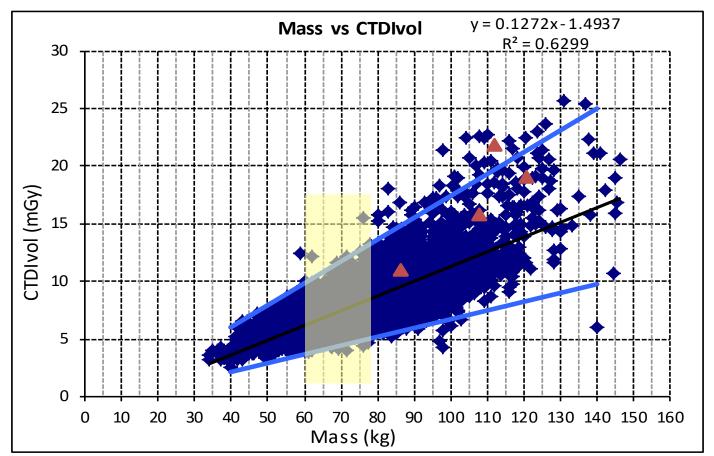
• Same mAs, same scan length



Absorbed dose to organ lower

Dose Audits - Patient size

- CTDIvol at 70 kg =~ 8 mGy
- Great uncertainty if take only a few data points from any weight
- If only a few data samples (even 20 30) standard weight more important



Courtesy E. Castellano, Royal Marsden, London

Dose Audits - Patient size Indicators

- Weight
- BMI (weight / (height x height)
- Lateral and AP dimensions, Effective diameter
- Professional judgement 'standard size', 'too large', 'too small' (Sutton BJR 2014, Palorini Eur Radiol 2014, Moorin JRP 2013)
- Water equivalent diameter (used in estimating SSDE) (IEC soon)



Paediatrics: body imaging : weight not age (EU RP185)

Dose Audits - Patient size Indicators

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Paediatrics: body imaging : weight not age (EU RP185)

Setting DRLs for a range of sizes ?

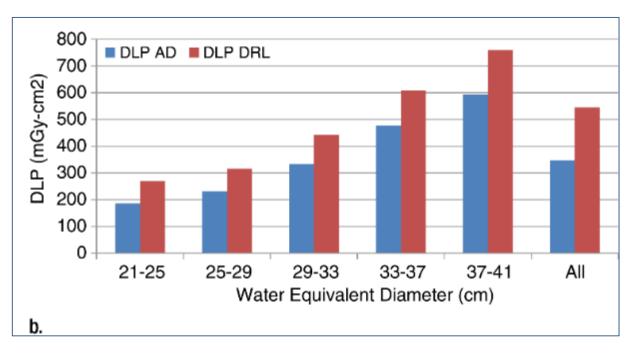
- Differences in the operation of tube current modulation systems affect the relationship between patient dose and size in different ways, so that translating tube current modulation settings in scanning protocols between CT scanners is not straightforward
- Relationships between the DRL quantities and patient size vary on different CT scanners
- Setting DRL values for different size ranges may be appropriate (manual methods not practical)
 ICRP 135

Size based DRLs

Figure 3. Graphs show abdomen and pelvis achievable doses (ADs) and diagnostic reference levels (DRLs). ..

(b) AD and DRL for abdomen and pelvis without contrast material—dose-length product (DLP





Original Research Medical Physics Free Access

U.S. Diagnostic Reference Levels and Achievable Doses for 10 Adult CT Examinations

Kalpana M. Kanal, Priscilla F. Butler ⊠, Debapriya Sengupta, Mythreyi Bhargavan-Chatfield, Laura P. Coombs, Richard L. Morin

What information to collect?

- How much of the scan protocol information should be collected?
 - kV, mA, scan time, recon algorithm, AEC
 - FBP or IR (and their parameters)
- Should it be ...
 - Just the exam name and dose index values ?

A compromise between too little information and too much – bearing in mind how you will process the information, and the people submitting data

ICRP 135: where information may give rise to key separation of system types this is important

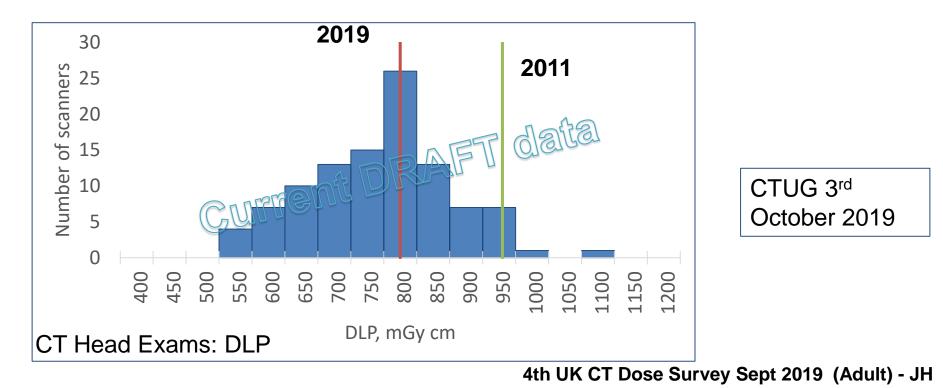
What information to collect? ICRP

- It is important that the data set in patient dose surveys for developing DRL values for CT includes:
 - detector technology
 - detector configuration
 - image reconstruction algorithm (FBP vs IR)
- So that differences between detector types and reconstruction algorithms are identified correctly.
- It may be useful to develop different DRL values locally for different CT technologies (e.g. single- vs multi-slice scanners, filtered back projection vs iterative reconstruction), even for the same procedure.

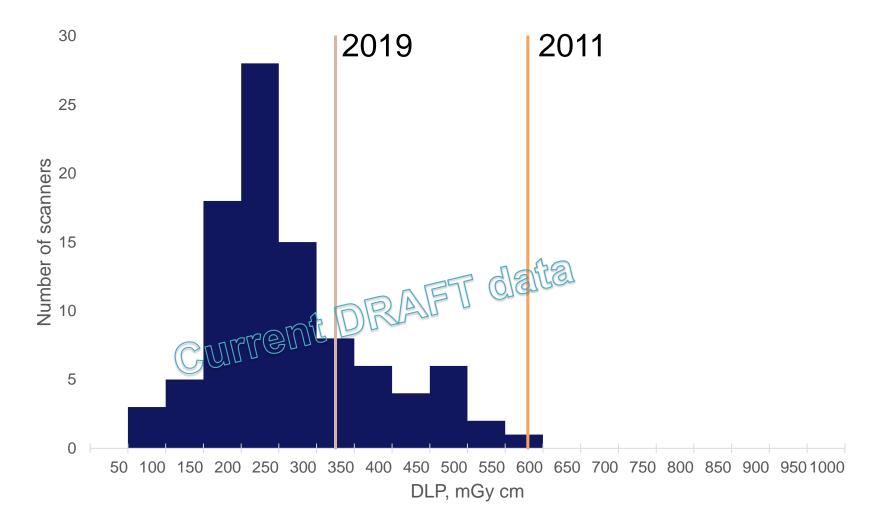
ICRP 135 (para. 214)

CT – Preliminary Results

- In general:
 - 10-30% reductions of proposed NDRL across the range of exams
 - >90% use AEC; 60 70% use IR



Chest exams: DLP



Separate dose by reconstruction technique

Examination	II	२	FE	3P	% Difference	
	CTDI _{vol}	DLP	CTDI _{vol}	DLP	CTDI _{vol}	DLP
Head	43.9	815	52.8	838	-17	-3
Paranasal sinuses	8.0	167	13.1	177	-39	-5
Cervical spine (C-spine)	15.2	431	22.0	492	-31	-12
Neck, chest, abdomen and pelvis	12.0	944	14.3	1060	-16	-11
Chest	8.8	290	10.7	374	-18	-22
Chest – high resolution	10.5	341	7.2	356	47	-4
Chest and abdomen	10.5	516	15.2	583	-31	-11
Chest-abdomen-pelvis (CAP)	11.1	734	14.6	754	-24	-3
CT pulmonary angiography (CTPA)	9.6	347	10.5	393	-8	-12
Abdomen and pelvis	12.8	640	14.0	670	-9	-5
Colonography/Virtual colonoscopy (VC)	6.0	842	8.0	835	-24	1
Kidney-ureters-bladder (KUB)	7.0	319	10.8	474	-35	-33
Urogram	9.4	974	9.2	966	3	1

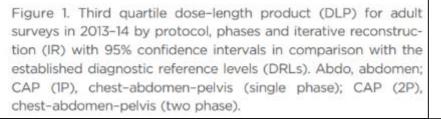
DRLs for new technology

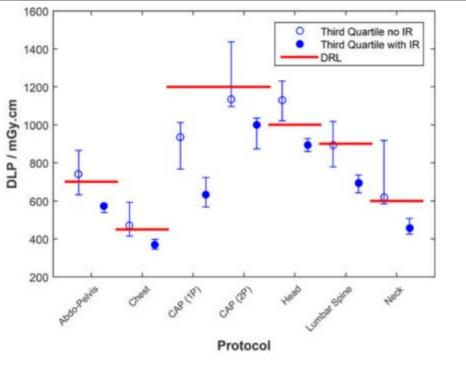
Evidence of dose saving in routine CT practice using iterative reconstruction derived from a national diagnostic reference level survey

P THOMAS, PhD, A HAYTON, MAppSc, BSc, T BEVERIDGE, PhD, P MARKS, BAppSc and A WALLACE, MAppSc, MSc

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Thomas P, Hayton A, Beveridge T, Marks P, Wallace A. Evidence of dose saving in routine CT practice using iterative reconstruction derived from a national diagnostic reference level survey. Br J Radiol 2015; 88: 20150380.

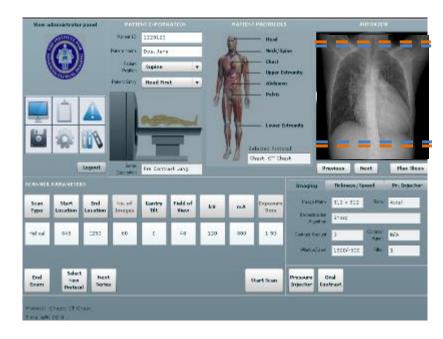


Key Questions we had

- Include the scan projection radiograph ?
- Contrast monitoring scans



SPR = scan projection radiograph = 'Scoutview', 'Topogram etc'

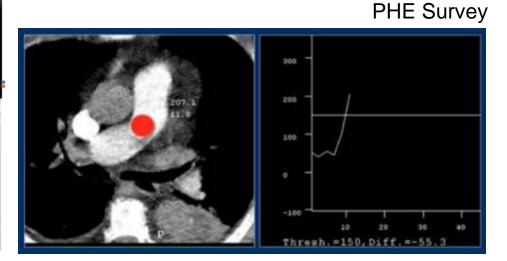




SPR and Bolus Tracking

- Exclude from the individual sequence data.
- Bolus tracking scans should be included in the total exam DLP
- SPRs may or may not be in total DLP (we ask if they are or aren't)





SPR = scan projection radiograph = 'Scoutview', 'Topogram etc'



Example Scan – Chest and Abdomen (Lung cancer)

• CCC_CHEST_ABDO_CONTRAST workflow:

	Sequence	CTDI	DLP	Exam
a.	Topogram (typical value)		7.96	
b.	Pre-contrast monitoring	1.16	1.20	
с.	Contrast monitoring (no IR)	1.16	1.20	
d.	Thorax CT(IR)	3.83	131.30	
e.	Abdomen CT (IR)	7.56	222.20	
	Exam	?	Total = 363.9	364.00 from scanner

- Need a consistent strategy as to how to quote CTDI for whole exam
- Should it be
 - 1. Exclude contrast and SPR, and give an average only of diagnostic image scans? Or
 - 2. Not quote CTDI for whole exam at all?
- PHE survey: Bolus tracking scans should be included in the total exam DLP
- PHE survey: SPR may or may not be given in total exam DLP (regardless it is only a small sue.edyvean@phe.gov.uk percent dose) (we want to know if they are or aren't)

CT Planning scans in Radiotherapy

Physics in Medicine & Biology





PAPER

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IPEM topical report: the first UK survey of dose indices from radiotherapy treatment planning computed tomography scans for adult patients

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Toshiba CTDI_{vol}

- For software version 4.63 or earlier, Toshiba scanners display maximum CTDI_{vol}, not average like all other vendors
 - *Typically* corresponds to scanners from before 2013
 - Scanners on later versions of software give average value
- For protocols that use the AEC system this will result in overestimation of the dose and may skew the national reference values for CTDI_{vol}

Does not affect DLP (based on average CTDI_{vol})

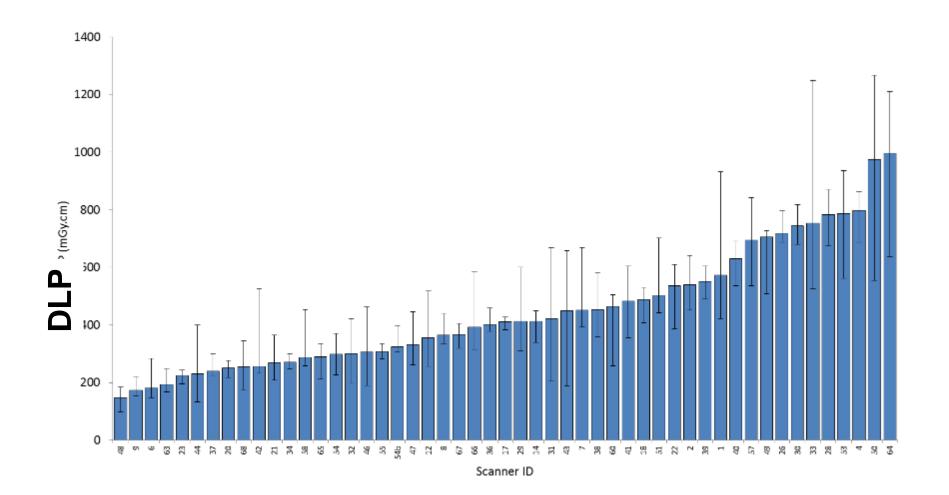
- All centres with Toshiba scanners installed prior to 2013 were asked to confirm the software version of their scanner
- If the data was from v4.63 or earlier;
 - The average CTDI_{vol} was excluding from the calculation of national reference values (DLP and scan length were left in)
 - CTDI_{vol} still included in plots for further discussion

(Tim Wood, Hull, UK. IPEM, CT in RT survey)



Institute of Physics and Engineering in Medicine

Lung 3D median DLP

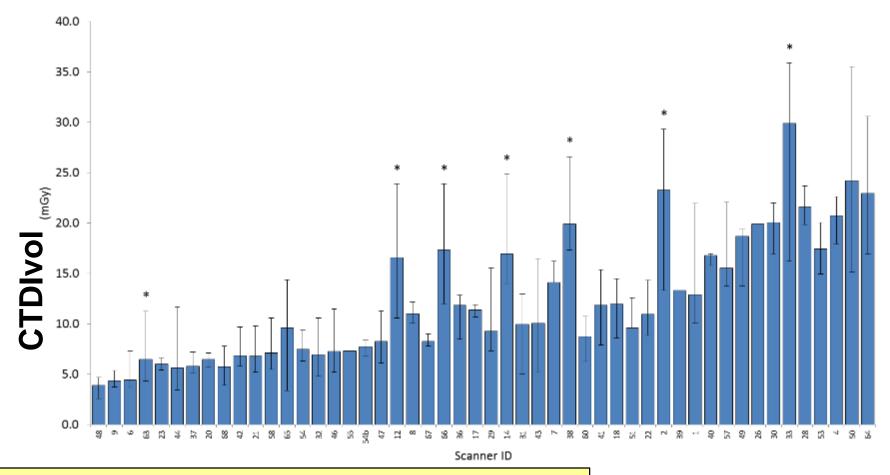


(Tim Wood, Hull, UK. IPEM, CT in RT survey)



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Lung 3D median CTDI_{vol}



* Indicates maximum CTDI on older Toshiba scanners using AEC

(Tim Wood, Hull, UK. IPEM, CT in RT survey)



Institute of Physics and Engineering in Medicine

High resolution chest CT

- Toshiba axial sequences appear to give CTDIw not CTDIvol
- 3 scanners with axial sequences, 1 mm beam width

Scanner	"CTDI _{vol} "	DLP
Aquilion CX	43	51
Aquilion One	50	60
Aquilion Prime	33	83

- Current CTDIvol NDRL is ~ 4 mGy
- The average CTDI_{vol} from other axial sequences in this study (n=11) is ~ 2 mGy
- Other manufactures appear to correct for step between scans, Toshiba do not



Dose Audits for DRLS

- Dose indicator (e.g. DAP,ESD or CTDI,DLP)
 - common examinations (e.g. chest CT) or high dose
 - Sample of standard size/weight patients
- Calculate the median^ value for each x-ray system, each exam



• ^UK previously used mean. UK currently ask for both: for retrospective comparison, and continue to future with median.

DRL: distribution of mean vs. median

DRL from distribution of

Examination	Mean	doses	Median	doses	% Difference		
	CTDI _{vol}	DLP	CTDI _{vol}	DLP	CTDI _{vol}	DLP	
Head	48.7	821	48.0	797	-1	-3	
Paranasal sinuses	12.0	173	11.6	165	-3	-4	
Cervical spine (C-spine)	17.6	473	17.6	443	0	-6	
Neck, chest, abdomen and pelvis	12.1	1026	10.0	904	-17	-12	
Chest	9.3	327	8.4	292	-10	-11	
Chest – high resolution	8.5	346	8.0	331	-5	-4	
Chest and abdomen	11.0	539	9.3	464	-15	-14	
Chest-abdomen-pelvis (CAP)	11.3	740	9.0	656	-20	-11	
CT pulmonary angiography (CTPA)	10.0	358	9.9	317	-2	-11	
Abdomen and pelvis	13.6	652	11.6	548	-15	-16	
Colonography/Virtual colonoscopy (VC)	7.2	857	6.8	820	-6	-4	
Kidney-ureters-bladder (KUB)	7.5	370	6.8	309	-10	-17	
Urogram	9.9	1010	8.9	913	-10	-10	

Note: this from well run dose audits. Errors may be greater for results of poorly run audits

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	3666 7 9111113	Median	7					
Add all	7+9+11+6+13+6+6+3+11 = 72 There are 9 numbers: 72 ÷ 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 out	liers			
Less robust for sk distributions		More robust for skewed distributions ghest value is 130 not 13:				
Nine numbe	ers: 7 9 11 6 <u>130 </u> 6 6 3 :	11				
Put in orde	r 3 666 7911	11 <u>130</u>	Mode	6		

Put in order	3666 7 91111 <u>130</u>	Median	7
Add all	7+9+11+6+ <u>130</u> +6+6+3+11 = 189 There are 9 numbers: 189 ÷ 9 = 8	Mean (average)	21

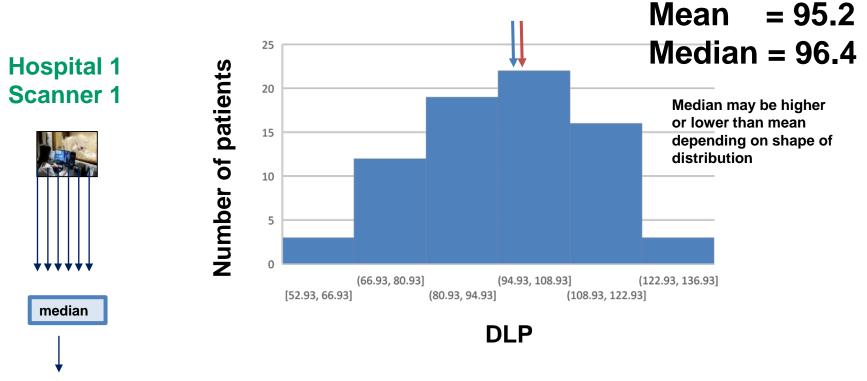


Mean versus Median

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	More robust for skewed distributions				
1.4 distributions – – n 1.2 same median – n 1.0- 0.8	node nedian neanNine numbers: 7 9 11 6 13 6 6 3 11Nine numbers: 7 9 11 6 13 6 6 3 11Mode60.25				
0.6 0.4 0.2 0.0	$\sigma = 1 \div 9 = 8 $ (average) 8				
0.0 0.2 0.4 0.6 0.8 1.0 1.2 1.4 1.6					

Public Distribution of data – mean and Median

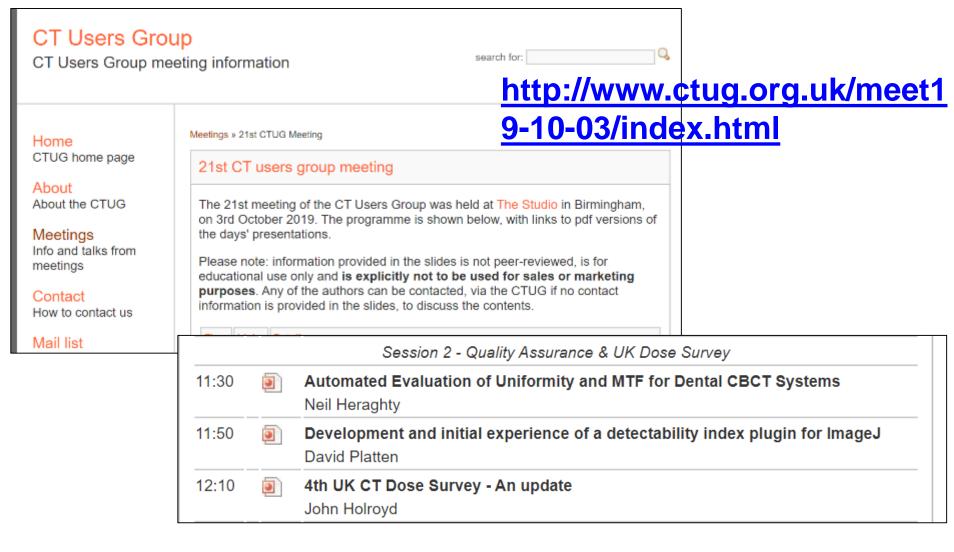
- 1. Distribution of one scanner's patient data / exam
 - Small data sample (standard weight)
 - Large data sample (no weights necessary if not available)



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Chest-abdo (lung cancer)

Update talk on 4th UK CT Survey



(Lots of talks on physics and CT : www.ctug.org.uk)

Same Spreadsheet - same generic info asked for Protocol details: scanner details

1	A	в	Ľ	D	E	F	Li Li	Н		J	K	L	M	N
1	1										4th	UK CT	Dose S	urvev
2	Public Health													
3 4	England													
4 5														
6	Body region (clinical indication)*: Abdomen and pelvis (Abscess)													
7												lease select	from drop	down list
8	If Other please give body region (clinical indication) details: domen and Pelvis (Nausea, Weight loss, abdo Pelvis Pain, Lethar													
9														
-	Hospital and		er Informa	tion									1	
-	Hospital Nam												xxxBridge	
2 Local system ID*: Diagnostic										-	r			
-	System manu												Siemens	
-	System mode			22.64.4	10 -t-l-								finition Edg	
-	Number of d			, 32, 64, 1	28, etc):							64 (128 sli		ing focal)
-	Year of manu Software ver		of scanner:									2018 Syngo CT VA84A		
-	Software ver	sion:										Syn	Igo CT VA84	A
8	Calibration	Data												
-	Error of indic		vol when la	ast checker	1 (+/-%)								6%	
ю И	Error or male		i i i i i i i i i i i i i i i i i i i	ist encener	a (., .,								070	
 22	Standard Pr	otocol S	ettings											
-	Local protoco											CCC_ABD	O_PELV_CC	NTRAST
4 Number of scan acquisitions* (e.g. 1 contrast & 1 non-contrast scan = 2 acquisitions):											1			
25														
26	Scanner/Pro	otocol Co	omments											
27	Please includ	le any oth	ner details a	and descri	ptions of y	our scan p	protocol							
8														
29														
0														

Protocol details: scout view details

Scout view details

Number of scout views:	
Does the total DLP (provided opposite) include the DLP from scout views?*	
Typical total DLP for all scout views (mGy.cm):	
Tube voltage (kV):	
Tube current (mA):	
Tuber current time (mAs):	
Imaged scan length (mm):	

Scout view details

Number of scout views:	1
Does the total DLP for each patient (below) include the DLP from scout views?*	No
Typical total DLP for all scout views (mGy.cm):	average 6.5
Tube voltage (kV):	120
Tube current (mA):	35
Tube current time (mAs):	
Imaged scan length (mm):	Average 480

Protocol details: scan details

Acquisition 1 details				See notes on s				
CTDI phantom size (cm) (i.e. 16 cr	n head or 32 cm body)*:		32 cm body	[a]				
Is Automatic Exposure Control (A	EC) used?*		No	[b]				
AEC name (e.g. AutomA, ZDOM, (CARE Dose 4D, SureExpose):			[c]				
AEC setting type (e.g. ref noise in	dex, reference mAs, etc):			[d]				
AEC setting value (e.g. SD 7.5, ref	mAs 200):		-	[e]				
minimum mA for AEC (where ap	plicable):		-	[f1]				
maximum mA for AEC (where ap	plicable):		-	[f1]				
mA where AEC is not used:		13	[f2]					
Is iterative reconstruction used?	ŧ		No					
Iterative recon type (e.g. ASIR, SA	AFIRE, iDose, AIDR):	-	[g]					
Iterative recon value (e.g. ASIR 40	0%, SAFIRE 3, iDose level 4):		-	[h]				
Radiation beam collimation	- Collimated beam width (mr	n):	10	[i]				
	- Number of slices:		1	63				
	- Detector size (mm) (e.g. 0.6	25,0.6):	10	[k]				
Is Automatic tube voltage selecti	on used? (eg. CarekV)		No					
If no, Fixed Tube voltage (kV):			120	[1]				
Tube rotation time (s):			1.5	[m]				
Primary image slice thickness (mi	m):		10	[n]				
Scan field of view (SFOV) (mm):				[0]				
Reconstruction field of view (DFC	Reconstruction field of view (DFOV) (mm):							
Axial or helical?	Axial	[9]						
Pitch (where applicable):			[1]					
Primary Reconstruction algorithm	n or kernel (e.g. B30; FC17; Sto	0	B30s	[5]				
Is contrast used?			None					
Anatomical landmarks for start a	nd end points	Start point (e.g. base of skull)	Carina	7				
		End point (e.g. vertex)	Carina	1				
	Comments		Pre-monitoring scan	1				

Protocol details: scan details

Split scan protocol settings (if applicable)

Acquisition 2 (if applicable)					
CTDI phantom size (cm) (i.e. 16 c	m head or 32 cm body)*:		32 cm body		
Is Automatic Exposure Control (A	AEC) used?*		No		
AEC name (e.g. AutomA, ZDOM,	CARE Dose 4D, SureExpose):		-		
AEC setting value (e.g. SD 7.5, re	f mAs 200):		-		
mA where AEC is not used:		13			
Is iterative reconstruction used?	\$		No		
Iterative recon value (e.g. ASIR 4					
Radiation beam collimation	m):	10			
	- Number of slices:		1		
	- Detector size (mm) (e.g. 0.	525,0.6):	10		
Is Automatic tube voltage select	ion used? (eg. CarekV)		No		
If no, Fixed Tube voltage (kV):			120		
Tube rotation time (s):			1.5		
Primary image slice thickness (m	ım):		10		
Scan field of view (SFOV) (mm):					
Reconstruction field of view (DF	OV) (mm):		300		
Axial or helical?		Axial			
Pitch (where applicable):					
Primary Reconstruction algorith	B30s				
Is contrast used?	IV				
Anatomical landmarks for start a	and end points	Start point (e.g. base of skull)	Carina		
		End point (e.g. vertex)	Carina		
	Comments		Contrast monitoring scans		

Protocol details: scan details

	comments		contrast monitoring scans							
Acquisition 3 (if applicable	0)									
			22 are bady							
CTDI phantom size (cm) (i.e. 1			32 cm body							
Is Automatic Exposure Contro			Yes CARE Dose4D							
	EC name (e.g. AutomA, ZDOM, CARE Dose 4D, SureExpose):									
	AEC setting value (e.g. SD 7.5, ref mAs 200):									
mA where AEC is not used:										
is iterative reconstruction use			Yes							
Iterative recon value (e.g. ASI		-	SAFFIRE strength 2							
Radiation beam collimation	 Collimated beam widt 	h (mm):	38.4							
	- Number of slices:		64							
	- Detector size (mm) (e.	g. 0.625,0.6):	0.6							
Is Automatic tube voltage sel	ection used? (eg. CarekV)		Yes							
If no, Fixed Tube voltage (kV):	:									
Tube rotation time (s):			0.5							
Primary <u>image</u> slice thickness	(mm):									
Scan field of view (SFOV) (mn	n):									
Reconstruction field of view (DFOV) (mm):									
Axial or helical?										
Pitch (where applicable):										
Primary Reconstruction algor	ithm or kernel (e.g. B30; FC1	7; Std)								
ls contrast used?										
Anatomical landmarks for sta	rt and end points	Start point (e.g. base of skull)								
		End point (e.g. vertex)								
	Comments		Thorax CT							

X PHE 4th UK Survey – patient data Public Health England

82.5	Essential fie	lds			
戀			4th UK CT Dose Survey	Acquisition 1 details	
Public Health	(blue)			CTDI phantom size (cm) (i.e. 16	cm head or 32 cm b
England	(101010)			Is Automatic Exposure Control	
-				AEC name (e.g. AutomA, ZDOM	
Body region (dinical indication	1*:			AEC setting type (e.g. ref noise	
- ,			Please select from drop down list	AEC setting value (e.g. SD 7.5, r	
If Other please give	e body region (clinical indicatio	n) details:		minimum mA for AEC (where a	
				maximum mA for AEC (where a	
Hospital and Scanner Informati	ion			mA where AEC is not used:	
Hospital Name*:				is iterative reconstruction used	2*
Local system ID*:				Iterative recon type (e.g. ASIR,	-
System manufacturer*:				Iterative recon value (e.g. ASIR	
System model*:				Rediation beam collimation	- Collimated beam
Number of detector rows (eg. 16, 32	, 64, 128, etc):			Addated again commuter	- Number of slices:
Year of manufacture of scanner:					- Detector size (mr
Software version:				Is Automatic tube voltage selec	tion used? (eg. Carel
				If no, Fixed Tube voltage (kV):	
Calibration Data				Tube rotation time (s):	
Error of Indicated CTDIvol when last	chacked (al. 56)			Primary image slice thickness (r	nm):
End of indicated croniol when ast	enecked (47-36)			Scan field of view (SFOV) (mm):	
				Reconstruction field of view (Di	FOV) (mm):
Standard Protocol Settings				Axial or helical?	
Local protocol name*:				Pitch (where applicable):	
Number of scan acquisitions* (e.g. 1	contrast & 1 non-contrast scan = 2 ac	quisitions):		Primary Reconstruction algorith	um or kernel (e.g. B3
				is contrast used?	

Acquisition 1 details										
CTDI phantom size (cm) (i.e. 16 cm head or 32 cm body)*:										
Is Automatic Exposure Control (A	VEC) used?*									
AEC name (e.g. AutomA, ZDOM,	CARE Dose 4D, SureExpose)									
AEC setting type (e.g. ref noise in	ndex, reference mAa, etc):									
AEC setting value (e.g. SD 7.5, re	f mAs 200):									
minimum mA for AEC (where app	plicable):									
maximum mA for AEC (where ap	plicable):									
mA where AEC is not used:	mA where AEC is not used:									
Is iterative reconstruction used?*										
Iterative recon type (e.g. ASIR, SAFIRE, iDose, AIDR):										
Iterative recon value (e.g. ASIR 40%, SAFIRE 3, IDose level 4):										
Radiation beam collimation	- Collimated beam width (mm):									
	- Number of slices:									
	- Detector size (mm) (e.g. 0.625,0.6):									
Is Automatic tube voltage selecti	ion used? (eg. CarekV)									
If no, Fixed Tube voltage (kV):										
Tube rotation time (s):										
	Primary image slice thickness (mm):									
Scen field of view (SFOV) (mm):										
Reconstruction field of view (DFOV) (mm):										
Axial or helical?										
Pitch (where applicable):										
Primary Reconstruction algorithm	n or kernel (e.g. B30; FC17; Std)									
Is contrast used?										

Patient det	tails						/This should	i ka ika						This should be th							
Patient No	Age (yrs)	t time of sc Weight (kg)	an: Height (cm)	Imaged	i langth (n Start position	End	distance from the star and.	rum the 4. to couch	tocol: Scan POV (mm)	стрі _{на} (mGy)*	DLP (mGy.cm)*	Imaged	Start position	distance from the centh start to co end.	e Hith	srotocol: Scan FOV (mm)	CTDI _{ve} [m@y]*	DLP (mSy.cm)*	can length (n Start position	nm) End position	Submit
1 2 3 4 5																					by patient
6 7 8 9 10																					(no ID info)
10 11 12 13 14																					

Patient details and dose

	Δ.	time of so				Acquisition 1									
	A	. time of st	.dn:	Scan	length (r	nm)	If differe	ent from prot	tocol:			Total DLP*			
Patient No	Age (yrs)	Weight (kg)	Height (cm)	Imaged length	Start position	End position	kV	CTDI phantom	Scan FOV (mm)	CTDI _{vol} (mGy)*	DLP (mGy.cm)*	(whole scan) mGy.cm			
1															
2															
3															
4															
5															

								Pre-monitoring					Cont	rast monitori	ng				
		t time of sc	201					Acquisition 1							A	cquisition 2			
	A	t time of sc	dn:	Scan	length (n	nm)	If different from protoc		If different from protocol:			Scan length (mm) If different from protocol:							
Patient No	Age (yrs)	Weight (kg)	Height (cm)	Imaged length	Start position	End position	kV	CTDI phantom	Scan FOV (mm)	CTDI _{vol} (mGy)*	DLP (mGy.cm)*	Imaged length	Start position	End position	kV	CTDI phantom	Scan FOV (mm)	CTDI _{vol} (mGy)*	DLP (mGy.cm)*
1							120	32 cm body		1.16	1.20				120	32 cm body		2.31	2.30
2							120	32 cm body		1.16	1.20				120	32 cm body		5.78	5.80
3							120	32 cm body		1.16	1.20				120	32 cm body		10.41	10.40
4							120	32 cm body		1.16	1.20				120	32 cm body		3.47	3.50
5							120	32 cm body		1.16	1.20				120	32 cm body		10.41	10.40
6							120	32 cm body		1.16	1.20				120	32 cm body		9.25	9.30
7							120	32 cm body		1.16	1.20				120	32 cm body		11.57	11.60
8							120	32 cm body		1.16	1.20				120	32 cm body		9.25	9.30
9							120	32 cm body		1.16	1.20				120	32 cm body		3.47	3.50
10							120	32 cm body		1.16	1.20				120	32 cm body		11.57	11.60
11							120	32 cm body		1.16	1.20				120	32 cm body		5.78	5.80
12							120	32 cm body		1.16	1.20				120	32 cm body		11.57	11.60
13							120	32 cm body		1.16	1.20				120	32 cm body		4.63	4.60
14							120	32 cm body		1.16	1.20				120	32 cm body		17.35	17.30
15							120	32 cm body		1.16	1.20				120	32 cm body		10.41	10.40
16							120	32 cm body		1.16	1.20				120	32 cm body		9.25	9.30
17							120	32 cm body		1.16	1.20				120	32 cm body		4.63	4.60
18							120	32 cm body		1.16	1.20				120	32 cm body		3.47	3.50

			The	orax CT				Abdomen-pelvis CT									
			Acqu	isition 3							Acquis	sition 4					
Sc	an length (m	m)	If dif	fferent from	protocol:			Scan length (mm)		Scan length (mm) If differe			otocol:			Total DLP*	
Imaged length	Start position	End position	kV	CTDI phantom	Scan FOV (mm)	CTDI _{vol} (mGy)*	DLP (mGy.cm)*	Imaged length	Start position	End position	kV	CTDI phantom	Scan FOV (mm)	CTDI _{vol} (mGy)*	DLP (mGy.cm)*	(whole scan) mGy.cm	Patient comments
			100	32 cm body		4.71	167.80				80	32 cm body		7.14	367.60	549.00	Large patient
			100	32 cm body		6.42	214.50				120	32 cm body		14.03	763.20	996.00	Large patient
			100	32 cm body		3.45	118.30				100	32 cm body		8.07	404.90	544.00	
			100	32 cm body		4.04	129.20				100	32 cm body		8.62	397.90	542.00	
			100	32 cm body		2.96	98.70				100	32 cm body		9.43	404.80	525.00	
			100	32 cm body		2.76	92.20				100	32 cm body		4.28	195.70	308.00	
			80	32 cm body		2.73	93.80				100	32 cm body		5.44	259.70	376.00	
			100	32 cm body		3.45	103.70				80	32 cm body		7.29	410.40	536.00	
			100	32 cm body		5.03	154.30				100	32 cm body		11.05	554.30	722.00	Large patient
			100	32 cm body		4.96	129.10				100	32 cm body		11.82	585.10	736.00	
			100	32 cm body		2.92	82.00				100	32 cm body		7.51	369.30	468.00	
			100	32 cm body		3.61	131.40				100	32 cm body		8.49	439.10	594.00	
			100	32 cm body		2.97	82.30				100	32 cm body		6.18	333.00	431.00	
			100	32 cm body		2.68	80.70				100	32 cm body		6.03	309.60	419.00	Large number of monitoring scans
			100	32 cm body		2.83	107.50				100	32 cm body		6.62	323.70	453.00	
			100	32 cm body		4.47	163.10				100	32 cm body		8.89	459.50	643.00	Large patient
			100	32 cm body		4.34	144.90				80	32 cm body		6.79	336.20	496.00	
			120	32 cm body		8.55	250.30				100	32 cm body		14.45	747.10	1010.00	Large patient
			100	32 cm body		3.53	141.20				100	32 cm body		5.91	305.50	469.00	
			100	32 cm body		6.55	191.00				100	32 cm body		14.74	762.50	967.00	Large patient
			100	32 cm body		5.32	147.60				120	32 cm body		17.02	906.50	1074.00	Large patient
			100	32 cm body		2.81	93.80				100	32 cm body		5.05	248.60	359.00	
			100	32 cm body		4.02	116.10				120	32 cm body		19.42	1038.30	1173.00	Large patient
			100	32 cm body		4.01	121.20				100	32 cm body		6.62	356.90	505.00	Large number of monitoring scans
			100	32 cm body		2.74	91.60				100	32 cm body		6.18	311.40	425.00	
			100	32 cm body		5.16	158.80				100	32 cm body		9.11	480.60	663.00	Large patient
			100	32 cm body		5.65	166.80				100	32 cm body		11.22	666.50	858.00	Large patient
			100	32 cm body		7.31	198.30				100	32 cm body		12.70	674.90	892.00	Large patient
			100	32 cm body		4.34	145.20				100	32 cm body		12.38	608.50	773.00	



Summary dose data from local audit

No of Patients	Mean Age at time of scan (yrs)	Mean Body Mass (kg)	Mean Total DLP* (whole scan)	Median Total DLP* (whole scan)	Comments on the data collection method (eg. inclusion criteria, data analysis method)

					Acquisition 1				
Mean CTDI _{vol} (mGy)*	Standard deviation	Median CTDI _{vol} (mGy)*	25th Percentile	75th Percentile	Mean DLP (mGy.cm)*	Standard deviation	Median DLP (mGy.cm)*	25th Percentile	75th Percentile

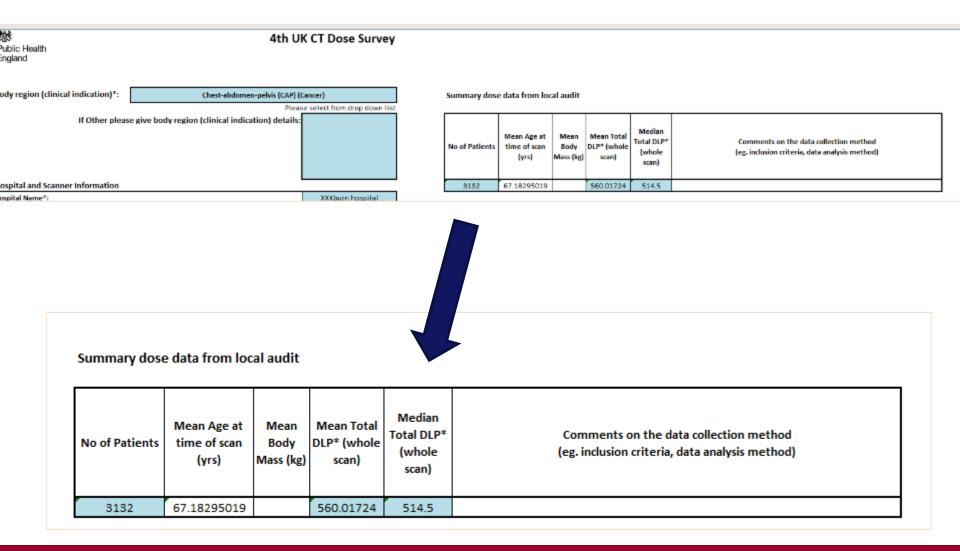
		_			Acquisition 2				
Mean CTDI _{vol} (mGy)*	Standard deviation	Median CTDI _{vol} (mGy)*	25th Percentile	75th Percentile	Mean DLP (mGy.cm)*	Standard deviation	Median DLP (mGy.cm)*	25th Percentile	75th Percentile

Or by summary data from local audit – for each system

Summary of Local audit - details and

If Other pleas	e IScout view details			
	Number of scout views:		1	٦
	Does the total DLP (provided opposi	te) include the DLP from scout views?*	Yes	
	Typical total DLP for all scout views (mGy.cm):	8.5	
	Tube voltage (kV):		120	1
Hospital and Scanner Information	Tube current (mA):		35	1
Hospital Name*:	Tuber current time (mAs):		NA	7
Local system ID*:	Imaged scan length (mm):		512	
System manufacturer*:	Acquisition 1 details CTDI phantom size (cm) (i.e. 16 cm h	ead or 32 cm bodv)*:	See notes on scanner spe 32 cm body	ecific he
System model*:	Is Automatic Exposure Control (AEC)		Yes	[b]
Number of detector rows (eg. 16, 32, 64			CareDose4D	[c]
Year of manufacture of scanner:	AEC setting type (e.g. ref noise index		Quality ref mAs	[d]
Software version:	AEC setting value (e.g. SD 7.5, ref m/	As 200):	180	[e]
	minimum mA for AEC (where applic	able):	NA	[f1]
Calibration Data	maximum mA for AEC (where applic	able):	NA	[f1]
Error of indicated CTDIvol when last chec	ke mA where AEC is not used:		NA	[f2]
	Is iterative reconstruction used?		Yes	
Standard Protocol Settings	Iterative recon type (e.g. ASIR, SAFIR	E, iDose, AIDR):	SAFIRE	[g]
Local protocol name*:	Iterative recon value (e.g. ASIR 40%,	SAFIRE 3, iDose level 4):	Strength 1	[h]
Number of scan acquisitions* (e.g. 1 cont	ras <u>Radiation beam</u> collimation	- Collimated Beam width (mm):	NA	[1]
		- Number of slices:	128	[]]
Scanner/Protocol Comments		- Detector size (mm) (e.g. 0.625,0.6):	0.6	[k]
-	Is Automatic tube voltage selection	used? (eg. CarekV)	Yes	
Please include any other details and des				[1]
	Tube rotation time (s):		0.5	[m]
	Primary <u>image</u> slice thickness (mm):		5	[n]
119 4th UK CT Dose 3	Su ^{Scan} field of view (SFOV) (mm):		NA	[0]
	Reconstruction field of view (DFOV)	(mm):	300	[p]
	Avial as halical?		the based	1 - 1

Summary of Local audit - details and doses



Summary of Local audit - details and doses

www. Public Health England 4th UK CT Dose Survey

ody region (clinical indication)*:	Chest-abdomen-pelvis (CAP) (C	Cancer)
	Pleas	e select from drop down list
If Other please	e give body region (clinical indication) details:	
ospital and Scanner Information		
ospital Name*:		XXXburn hospital
ocal system ID*:		RBC1
/stem manufacturer*:		Siemens
/stem model*:		Definition AS
umber of detector rows (eg. 16, 32, 64,	128, etc):	128

Summary dose data from local audit	Summary	dose	data	from	local	audit	
------------------------------------	---------	------	------	------	-------	-------	--

No of Patients	Mean Age at time of scan (yrs)	Mean Body Mass (kg)	Mean Total DLP* (whole scan)	Median Total DLP* (whole scan)			n the data collectio riteria, data analys		
3132	67.18295019		560.01724	514.5					
	_								
					Acquisition 1				
Mean C (mG	Standard deviation	Median CTDI _{vol} (mGy)*	25th Percentile	75th Percentile	Mean DLP (mGy.cm)*	Standard deviation	Median DLP (mGy.cm)*	25th Percentile	75th Percentile

Generally only get total Exam DLP data

Summary dose data from local audit

P	No of Patients	Mean Age at time of scan (yrs)	Mean Body Mass (kg)	Mean Total DLP* (whole scan)	Median Total DLP* (whole scan)	Comments on the data collection method (eg. inclusion criteria, data analysis method)
	3132	67.18295019		560.01724	514.5	

Teaching material

- Basic CT
 - www.impactscan.org
- Physics UK Group
 - www.ctug.org.uk
- CTISUS.org







Protecting and improving the nation's health

DRLs and exposure monitoring in CT:

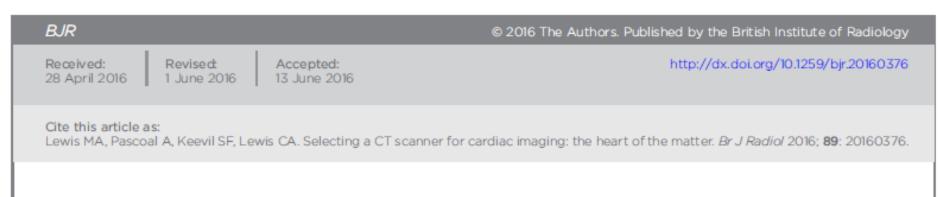
quantities, procedures, methods, international experience

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

Reports on Cardiac CT



REVIEW ARTICLE

Selecting a CT scanner for cardiac imaging: the heart of the matter

¹MARIA A LEWIS, Msc, ^{2,3}ANA PASCOAL, PhD, MIPEM, ^{1,2,4}STEPHEN F KEEVIL, PhD, FIPEM and ^{2,3}CORNELIUS A LEWIS, PhD, FIPEM

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Address correspondence to: Mrs Maria Anna Lewis E-mail: maria_anna_lewis@yahoo.co.uk

UK Cardiac CT Course, S Edyvean 2017

Reports on Cardiac CT

NICE National Institute for	NICE	NICE	Standards	Evidence	Sign in
Health and Care Excellence	Pathways	Guidance	and indicators	services	
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New generation cardiac CT scanners (Aquilion ONE, Brilliance iCT, Discovery CT750 HD and Somatom Definition Flash) for cardiac imaging in people with suspected or known coronary artery disease in whom imaging is difficult with earlier generation CT scanners

Diagnostics guidance [DG3] Published date: January 2012

Cardiac CT

- Cardiac CT BIR webinar 9 May 2016 (http://www.bir.org.uk/webinars-on-demand)
- Market review: Advanced CT scanners for coronary angiography CEP10043, March 2010

http://www.impactscan.org/reports/CEP10043.htm

Purchasing and Supply Agency

Centre for Evidence-based Purchasing

Advanced CT scanners for coronary angiography. CEP10043, Mar-10

This market review is intended to help prospective purchasers make informed choices and achieve best value from investment in high-end CT systems for Coronary CT Angiography applications. It should be read in conjunction with CEP's buyer's guide to multi-slice CT scanners (CEP08007) and the associated comparative specification reports (CEP08027, CEP08028).

Electronic access to a version of this report is available from the CEP website.

Advanced CT acc	nnoro for opronory
angiography	nners for coronary
CEP10043	
March 2010	