Dose assessment and optimization of acquisition protocols for CT simulation

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Joint ICTP-IAEA Workshop on Radiation Protection in Image-Guided Radiotherapy (IGRT)

Summary

Dose assessment

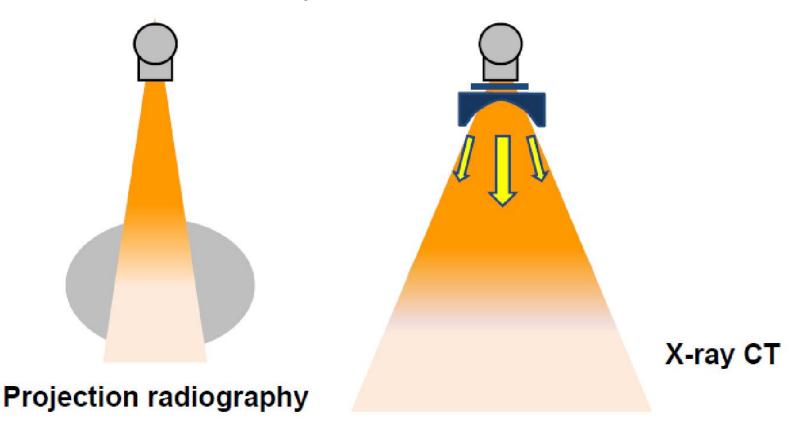
- Dose distribution from a CT scan
- Dose Metrics used in CT
 - CTDI (CTDI∞CTDI100, CTDIair, CTDIw, CTDIvol)
 - DLP
 - Effective dose
 - CTDIvol Wide beam
 - SSDE (size specific dose estimate)
- CTDI infinity

DRLs

Dose Optimization

Dose distribution in Scan Plane

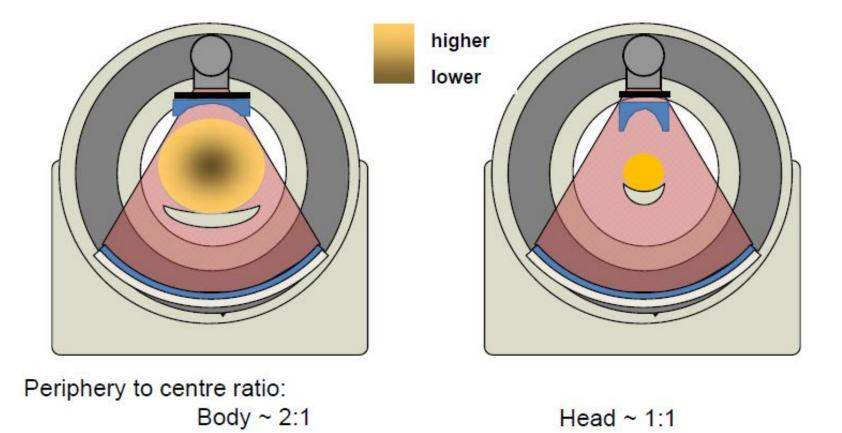
In PR part of the body irradiated



In CT whole body irradiated

BIR CT Optimisation June 2014 S Edyvean

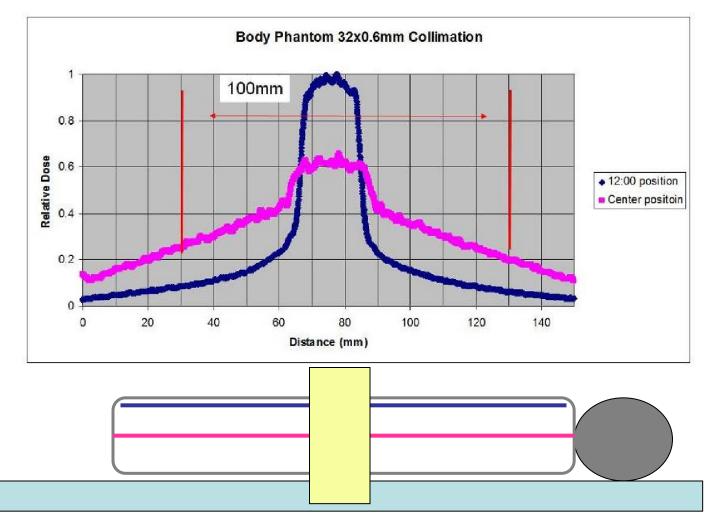
Dose distribution in Scan Plane



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Dose distribution in CT

Dose profile from one slice

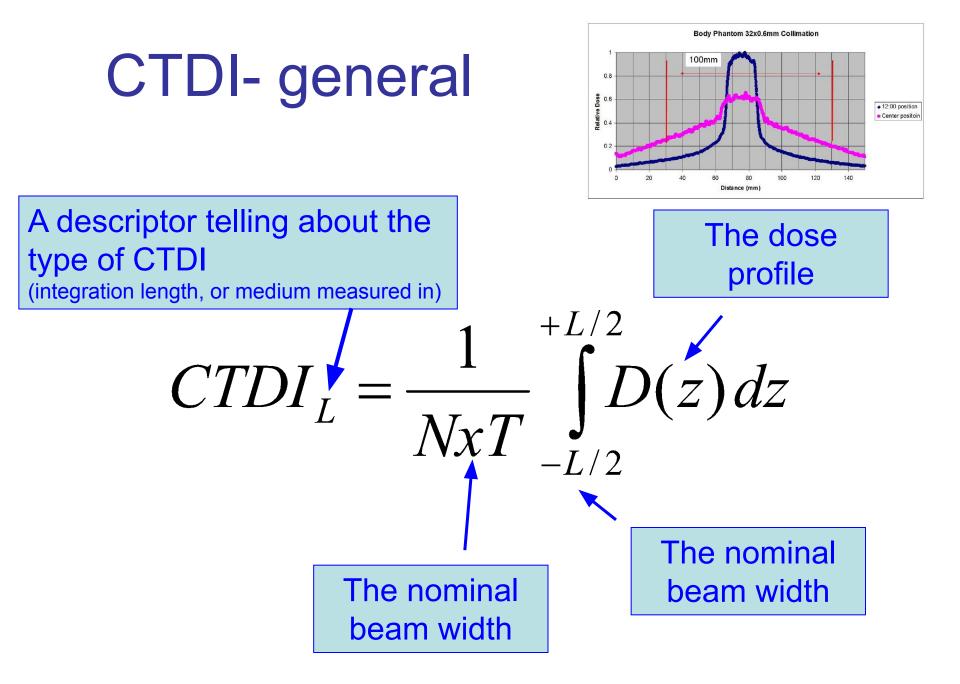


Dose metrics in CT

- MSAD Multiple Scan Average Dose
- CTDI Computed Tomography Dose Index*
- DLP Dose Length Product
- E Effective Dose
- SSDE Size Specific Dose Estimate

CTDIair CTDI100 CTDIw CTDIvol CTDI∞

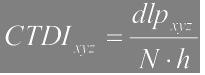
*



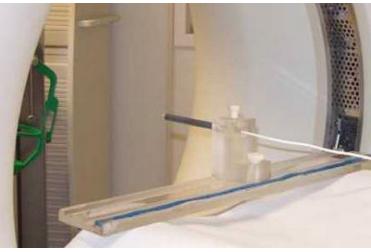
CTDI_{air}

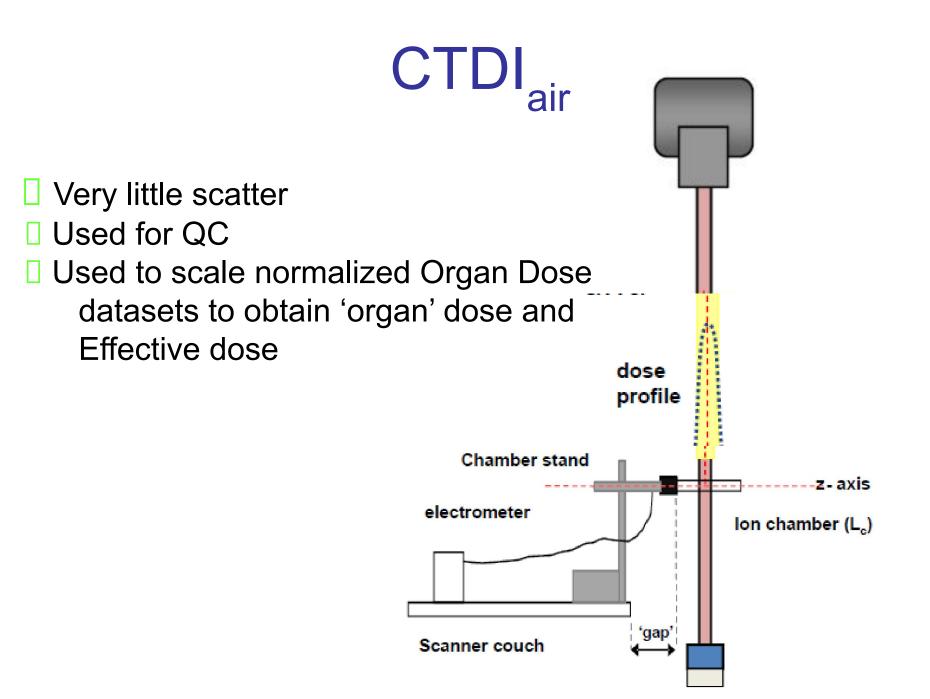
CTDIair

- Standard 100-mm pencil chamber dosimeter, must be placed in air at the isocenter of the scanner.
- The measure is expressed in terms of the DLP for a single slice, mGy*cm
- To obtain the CTDIair It is necessary do divide by the nominal thickness of the sliq



- N = n° of slices acquired simultaneously
- h = Slice thickness in cm





CTDI_{air}

Accuracy of set up

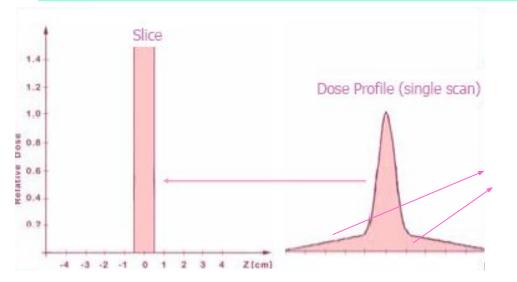
- iso-centre : +/- 10 mm, 0.1% error
- along chamber: +/- 10 mm, 0.3 % error
- tilt or twist: < 5 degrees, 0.1% error

Changes in CTDIair due to:

- Tube output
- Focal spot
- Beam width collimation
- Beam filtration

CTDI_{air} is one of the most sensitive and valuable parameters to measure

CTDI₁₀₀



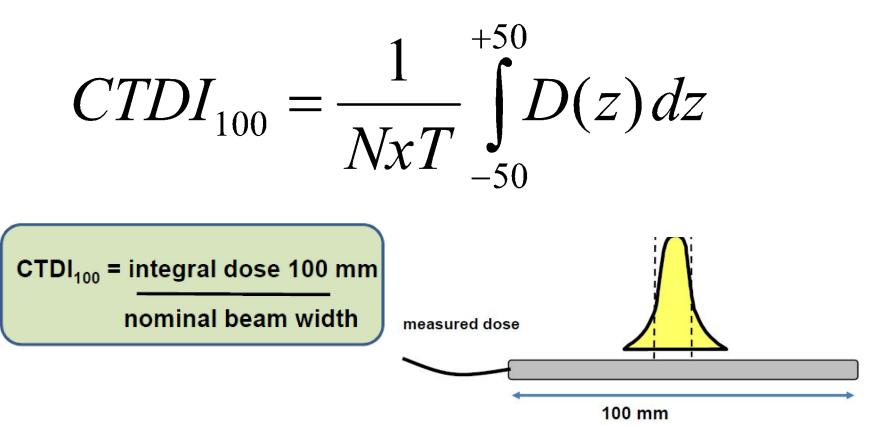
The queues before and after the slice thickness must be added to the dose in the slice thickness

The CTDI (CT Dose Index) is the standard dose D (z), imparted by a single axial acquisition to a standard 100-mm pencil chamber dosimeter inside a PMMA phantom along a line parallel to the axis of rotation, divided by the nominal thickness of the beam:

$$CTDI_{100} = \frac{1}{nh} \cdot \int_{-50}^{50} D(z) dz$$

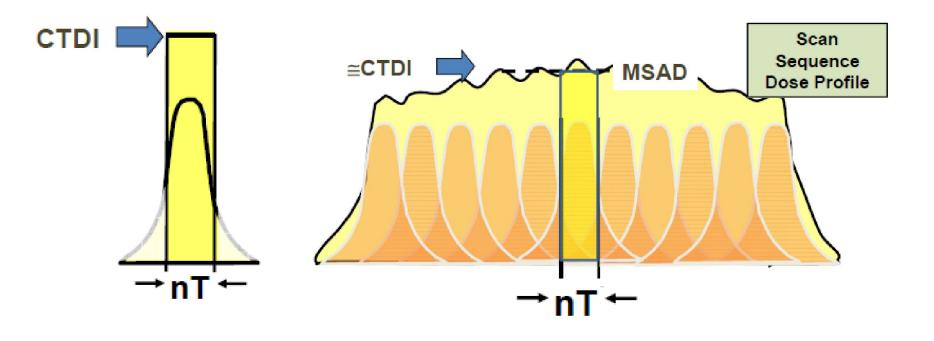
CTDI₁₀₀

- 100 mm long ion chamber used
- One 'slice' scanned
- The dose from the profile is collected over 100 mm
 That value is divided by the nominal beam width



CTDI₁₀₀

Is a calculation from a single slice measurement
 It represents the average dose to the centre of a scanned length (100 mm)



How accurate is the manufacturer displayed $CTDI_{100}$?

Variability of the discrepancy between manufacturer and measured $CTDI_{100}$ values by scanner type, acquisition parameters and phantom size

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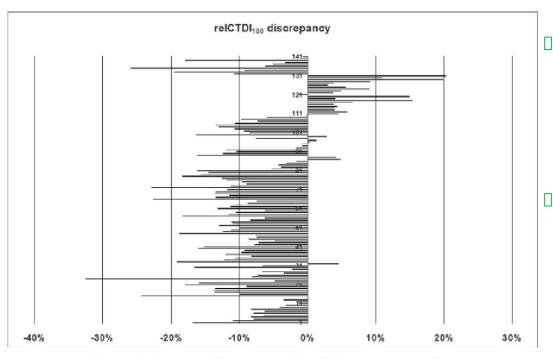


Fig. 1. Relative CTDI100 discrepancy (%) for all the measurements taken.

- The inaccuracies in the CTDI100 can lead to higher than 20% inaccuracies in the displayed CTDIvol, which is the suspension level indicated in the EC RP N.162.
- There is need for individual calibration of every single X-ray tube in CT by the manufacturers and the necessity of including this check in the quality control programs for CT equipment.

How accurate is the manufacturer displayed $CTDI_{100}$?

National reference levels of CT procedures dedicated for treatment planning in radiation oncology

Ana Božanić ^{a, b, *}, Doris Šegota ^a, Dea Dundara Debeljuh ^{a, b, c}, Manda Švabić Kolacio ^a, Đeni Smilović Radojčić ^{a, b}, Katarina Ružić ^d, Mirjana Budanec ^e, Mladen Kasabašić ^f, Darijo Hrepić ^g, Petra Valković Zujić ^{h, i}, Marco Brambilla ^j, Mannudeep K. Kalra ^{k, l}, Slaven Jurković ^{a, b}

| Center | Scout accuracy Δ/mm | ∆CTDI _{vol} (16 cm phantom)/% | ∆CTDI _{vol} (32 cm phantom)/% |
|--------|---------------------------|---|---|
| | 0 | -14.4 | -9.4 |
| 2 | 0 | 12.0 | 14.7 |
| 3 | 0 | 6.2 | 9.1 |
| 1 | 0 | -19.5 | -16.7 |
| 5 | 0 | -4.7 | -1.6 |
| 5 | 0 | -7.1 | -10.5 |
| 7 | 0 | 7.3 | 8.9 |
| 3 | 0 | 11.7 | -0.6 |

| Table 1 | |
|--|--|
| Quality control results for CTDIvol verification and scout accuracy. | |

Weighted CTDI (CTDI_w)

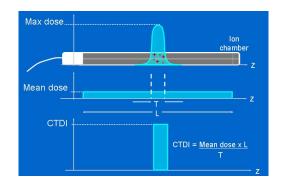
The dose distribution imparted by a CT scan is much more homogeneous than that imparted by radiography, but is still somewhat larger near the skin than in the centre of the body. The *weighted* CTDI was introduced to account for this:

They are obtained by placing the pencil dosimeter inside two PMMA cylindrical phantoms of 32 and 16 cm diameter representing an adult Body and a Head

$$CTDI_{w} = \frac{1}{3} \cdot CTDI_{100,C} + \frac{2}{3} \cdot \langle CTDI_{100,p} \rangle$$

Dividing by the mAs we obtained the normalized

nCTDIw



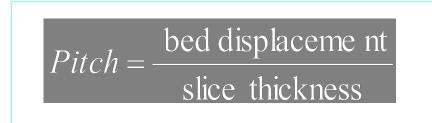


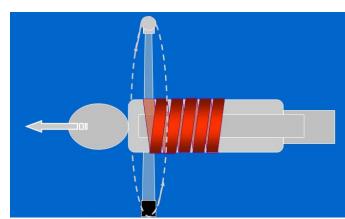
Must be shown in the console

Represents the average dose in scan plane of a 100 mm scan

Volume CTDI (CTDI_{Vol})

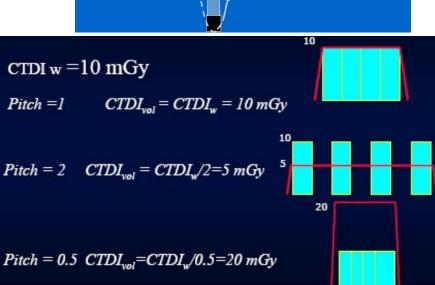
In the Spiral CT during a complete rotation of the Xray tube –detectors assembly we have a movement of the bed along the Z axis;





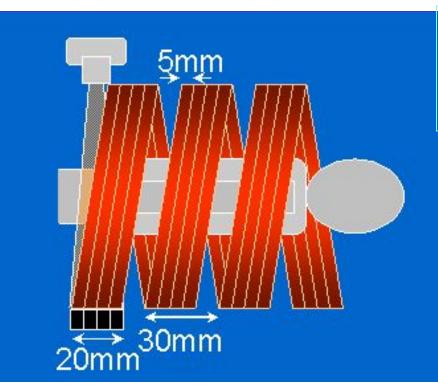
Dividing the nCTDIw by the pitch we obtain:





CTDI_{Vol} in MDCT

<u>MDCT</u>



$$Pitch = \frac{bed displacement}{primary collimatio n}$$

Primary collimation = n° acquired slices * slice Thickness

p=30/20 =1.5

CTDI_{vol}

Measured at acceptance and after major changes in CT scanner (Tube replacement)

Used to establish dose reference levels

Not to be used in QC

CTDIvol (and DLP) displayed on console

Dosimetric Indexes in CT: Factors P_B and P_H

The relation between *CTDI*w and *CTDI*air depends on the scanner type used for the examination and on the dosimetry phantom considered.

For the purpose of dose estimation, the ratio of both quantities is defined for the standard head (H) and body (B) CT dosimetry phantom

$$P_H = \frac{CTDI_{w,H}}{CTDI_{air}}$$
 and $P_B = \frac{CTDI_{w,B}}{CTDI_{air}}$,

Dosimetric Indexes in CT: Factors PB and PH

Table 1 Summary of characteristic performance parameters^a for four single-slice and six multi-slice CT systems used for dose calculation in this study

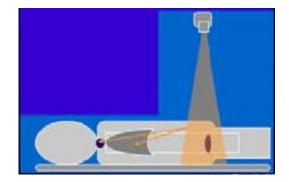
| Manufacturer | Scanner | Abbr. | N | U _{ref} (kV) | h _{ref} (mm) | dz (mm) | Head mode | | | Body mode | | | |
|------------------|-----------------|-------|----|--------------------------|--------------------------|------------|-----------|----------------|-----------------|-----------------------------------|----------------|-----------------|--|
| | | | | | | | (mGy/mAs) | P _H | k _{CT} | nCTDI _{w,B} (mGy/mAs) | P _B | k _{CT} | |
| General Electric | LX/i | G-1 | 1 | 120 | 10 | 0 | 0.152 | 0.66 | 0.80 | 0.072 | 0.31 | 0.65 | |
| Philips | Tomoscan AV | P-1 | 1 | 120 | 10 | 0 | 0.150 | 0.75 | 0.90 | 0.080 | 0.40 | 0.80 | |
| Siemens | Somatom Plus 4 | S-1 | 1 | 120 | 10 | 0 | 0.146 | 0.82 | 1.00 | 0.083 | 0.47 | 1.00 | |
| Toshiba | XVision | T-1 | 1 | 120 | 10 | 0 | 0.162 | 0.63 | 0.80 | 0.065 | 0.30 | 0.65 | |
| General Electric | Lightspeed QX/i | G-4 | 4 | 120 | 5 | 3.0/4.0b | 0.182 | 0.64 | 0.80 | 0.094 | 0.39 | 0.80 | |
| Philips | Mx8000 Quad | P-4 | 4 | 120 | 5 | 1.7 | 0.130 | 0.75 | 0.90 | 0.067 | 0.39 | 0.80 | |
| Siemens | Volume Zoom | S-4 | 4 | 120 | 5 | 1.7 | 0.200 | 0.76 | 0.90 | 0.083 | 0.49 | 1.00 | |
| Toshiba | Aquilion | T-4 | 4 | 120 | 8 | 3.0 | 0.189 | 0.67 | 0.80 | 0.107 | 0.30 | 0.65 | |
| Philips | Mx8000 IDT | P-16 | 16 | 120 | 1.5 | 3.0 | 0.130 | 0.75 | 0.90 | 0.067 | 0.39 | 0.80 | |
| Siemens | Sensation 16 | S-16 | 16 | 120 | 1.5 | 3.0 | 0.190 | 0.76 | 0.90 | 0.070 | 0.41 | 0.80 | |

^a Definition of scanner parameters: N, number of simultaneously acquired slices; U_{ref} , reference voltage for ${}_{n}CTDI_{w,H/B}$; h_{ref} , slice collimation for ${}_{n}CTDI_{w,H/B}$; dz, width of penumbra; ${}_{n}CTDI_{w,H/B}$, normalized $CTDI_w$ for head or body mode; $P_{B/H}$, phantom factor for head or body mode; k_{CT} , scanner-specific correction factor. ^b Value depends on focal spot size.

Dose Lenght Product DLP

By doubling the scan lenght, the CTDI_{vol} doubles? NO

The CTDI indexes are dose indicators. As such they refer to the dose in a specific point. They do no take into account the length of the acquisition.



The product of the CTDI times the length of the acquisition is the Dose Length Product :

$$DLP = _{n} CTDI_{w} \cdot mAs \cdot n \cdot h$$

The DLP can be used to estimate the Effective Dose

It is an index of the global irradiation of the patient.

Effective Dose ED

To reflect the combined detriment from stochastic effects due to the equivalent doses in all the organs and tissues of the body, the equivalent dose in each organ and tissue is multiplied by a tissue weighting factor, W_T , and the results are summed over the whole body to give the effective dose E

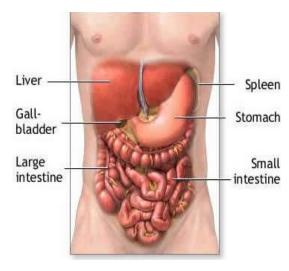
$$E = \sum_{\mathrm{T}} w_{\mathrm{T}} \sum_{\mathrm{R}} w_{\mathrm{R}} D_{\mathrm{T,R}} \quad \text{or} \quad E = \sum_{\mathrm{T}} w_{\mathrm{T}} H_{\mathrm{T}}$$

where HT or $w_R D_{T,R}$ is the equivalent dose in a tissue or organ, T, and w_T is the tissue weighting factor. The unit for the effective dose is the same as for absorbed dose, J kg⁻¹, and its special name is Sievert (Sv).

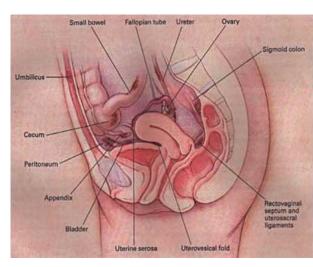
Effective Dose ED

ABDOMEN ABDOMEN PELVIS

PELVIS



| | | | $W_{T}H_{T}$ |
|--------------|----------------|----------------|--------------|
| Organs | W _T | H _T | |
| Gonads | 0,08 | 31,385 | 2,511 |
| Bone Marrow | 0,12 | 12,329 | 1,479 |
| Colon | 0,12 | 23,233 | 2,788 |
| Lung | 0,12 | 7,359 | 0,883 |
| Stomach | 0,12 | 24,794 | 2,975 |
| Bladder | 0,04 | 32,403 | 1,296 |
| Breast | 0,12 | 1,355 | 0,163 |
| Liver | 0,04 | 22,636 | 0,905 |
| Oesophagus | 0,04 | 1,063 | 0,043 |
| Thyroid | 0,04 | 0,097 | 0,004 |
| Skin | 0,01 | 8,826 | 0,088 |
| Bone Surface | 0,01 | 16,392 | 0,164 |
| Remainder 2 | 1,456 | | |
| Effective | 14,8 | | |



Tissue weighting factors (W_{T}) derived from whole population

What the scanner shows

| Name: | PHYSICS T | EST | ID:PF | IY12345 | Protoco | ol:10.4 | | | | | | | | | | | | |
|---------------------------------|-----------------------------|-------------------|--|------------------|-------------------------|------------------|------|----------------|-----|---------------|-------|-----|----------------|---|---------------|---|---------|---|
| Anatomi | | | | | mical XY | Dose Information | | | | | | | | | | | | |
| | | | Patient Orie Head First Patient Posi | | Ir | Images | | CTDIvol mGy | | DLP mGy∙cm | | | Dose Eff. % | | Phantom cm | | | |
| Series Description | | | | e | | 1-5 25.77 | | 134.67 | | | 89.31 | | Body 32 | 2 | | | | |
| Add | Spli | t De | lete B | iopsy | Smart | Pn | | 6-10 | | 25.7 | 7 | 134 | .67 | | 89.31 | | Body 32 | 2 |
| Group Current Selected Group Rx | | Rx | Prep Rx mA | | 1 | 11-15 | | 25.7 | 7 | 134 | .67 | | 89.31 | | Body 32 | 2 | | |
| Images | Scan Type | Start Location | End | No. of Images | Thick Speed | C _{im} | | | | | | | | | | | | |
| 1-5 | Helical Full 0.6 sec. | \$15.750 | 14.250 | 5 | 5.0 27.50 1.375:1 | 5.000 | | | | | | | | | | | 1.5 | |
| 6-10 | Helical Full 0.6 sec. | \$15.750 | 14.250 | 5 | 5.0 27.50 1.375:1 | 5.000 | 50.0 | Large Body | 120 | 650 11.57~ | 1.1 | 3.3 | 1.3 | N | N | N | 2.0 | |
| | Helical | | | | 5.0 | | | | | | | | | | | 1 | | |

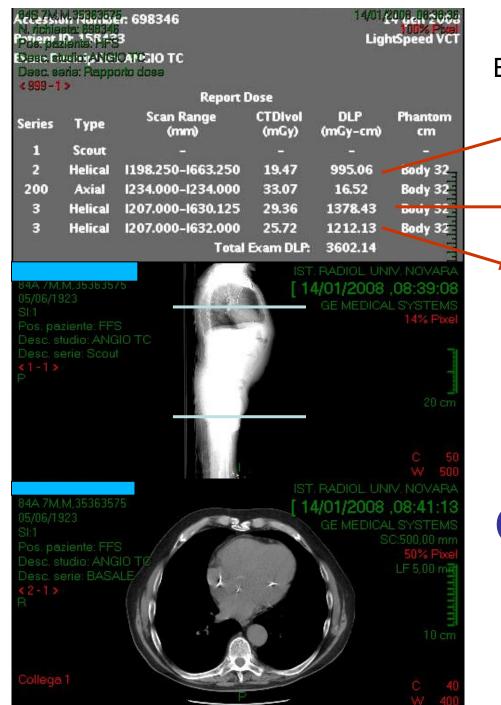
GE Scanner

ED_{DLP} Conversion Factors

E = DLP x k k values region specific

Guidelines EUR16262EN

| Region of body | Normalized Effective Dose E _{DLP} (mSv mGy ⁻¹ cm ⁻¹) |
|----------------|--|
| Head | 0.0023 |
| Neck | 0.0054 |
| Chest | 0.017 |
| Abdomen | 0.015 |
| Pelvis | 0.019 |



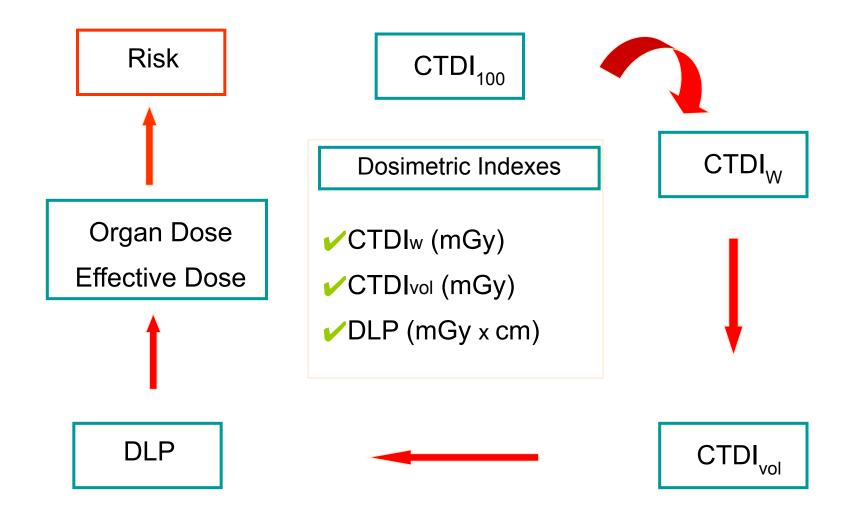
 $E_{DLP} (mSv mGy^{-1} cm^{-1}) = 0.017 - 0.019$ E (mSv) = 0.017 x 995.06 = 16.9 E (mSv) = 0.017 x 1378.43 = 23.4 E (mSv) = 0.019 x 1212.33 = 23.0 $E_{Tot} (mSv) = 63.3$

CT Abdomen- Pelvis Multiphase

CT Head

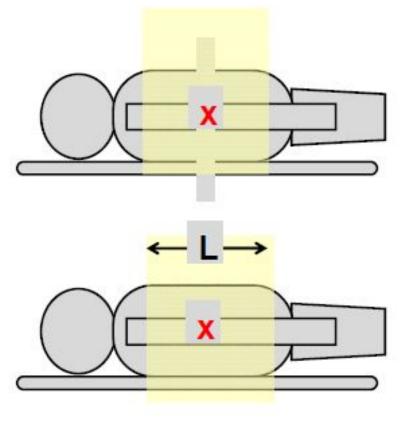
| 78A 4M,F,100116381 25/12/1931 SI:2 ID richiesta: 1256145 Pos. paziente: HFS Desc. serie: Dose Info | Fime: Acc.Number: Fotal DLP: BRALE | Mag 17, 2010, 15:42:59 1256145 766.0 mGy*cm | SCHED.MED [17/05/2010 ,15:42:59] 120kV SC:500,00 mm |
|--|---|--|--|
| < 80140-2> | Dose # Description 1 2 3mm | Mode CTDI DLP [mGy] [mGy*cm] Surview 0.0 0.00 Surview 0.0 0.00 Axial 53.2 766.02 | |
| 78A 4M,F,100116381 25/12/1931 SI:1 Pos. paziente: HFS Desc. studio: 12088-TC Desc. serie: 1.5mm OSSO < 3-11.5mm OSSO > | MaggioredellaCo SCHED.M [17/05/2010 ,15:43 Philips Brillianci Sc500.00 54% F LF 1,50 | EDLP (mSv mGy ⁻¹ c | m ⁻¹) = 0.0023 |
| R Collega 1 | | E (mSv)= 0.0023 | x 766.02 = 1.76 |

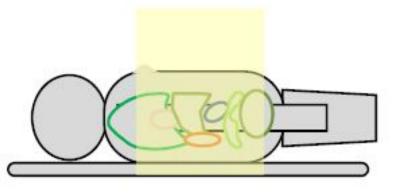
Relationship between Dosimetric Indexes in CT



Conclusions ?

- CTDIvol (mGy)
 - ~ represents local absorbed dose
 - DLP = CTDIvol x L (mGy.cm)
 - represents total absorbed dose
 - $-\sim$ represents relative risk
 - Effective dose (mSv)
 - Sensitivity of organs accounted for
 - ED = DLP x k
 - k values region specific





CTDI is not patient dose



Good for QC

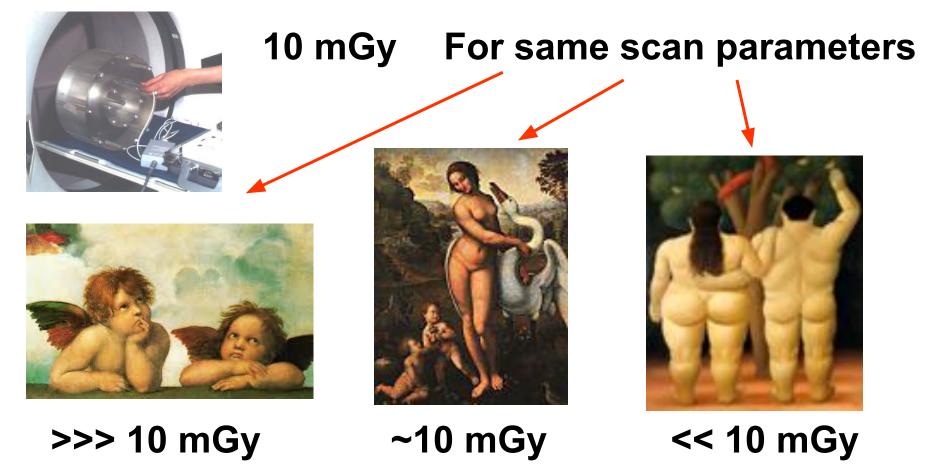
 A good indicator of relative dose between protocols, scanners and standard size patients



- Patients are not 16 or 32 cm cylinders of Perspex
- The integration length (≡ scan length) is not 100 mm
- Patients come in different sizes

Patients come in different sizes

- Patients come in different sizes
- But we quote CTDIvol to the same size phantoms



AAPM Report No. 204



Size-Specific Dose Estimates (SSDE) in Pediatric and Adult Body CT Examinations

Report of AAPM Task Group 204, developed in collaboration with the International Commission on Radiation Units and Measurements (ICRU) and the Image Gently campaign of the Alliance for Radiation Safety in Pediatric Imaging





Size-specific Dose Estimation for CT: How Should It Be Used and What Does It Mean?¹

James A. Brink, MD Richard L. Morin, PhD

wing to rising concerns about a metric of radiation output, not of paonizing radiation from medical tient dose. The exposure to radiation is maging, the National Council the same whether measured in a block

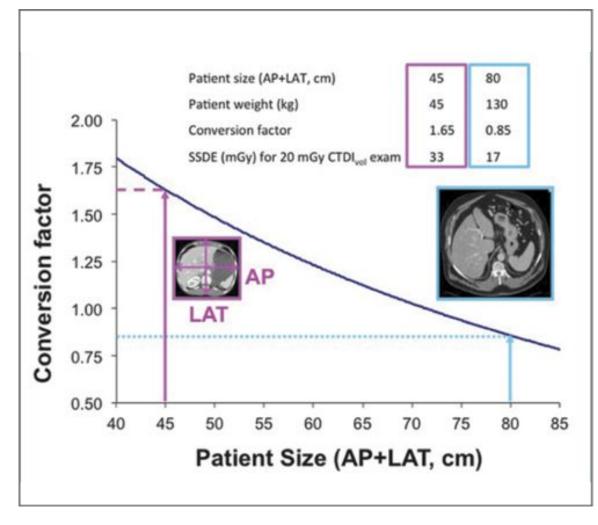
December 2012 Radiology, 265, 666-668.

This value, given in mGy, is an adjusted CTDI value based on the patient's size
 Uses Effective diameters

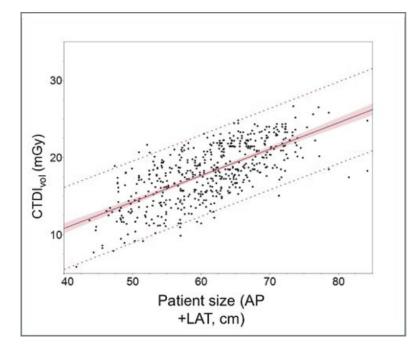
Eff Diameter =
$$\sqrt{APxLAT}$$

Small BodiesLarge Bodies

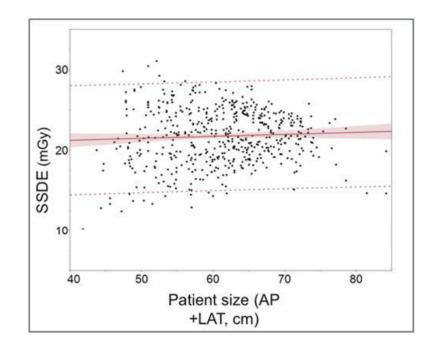




Demonstration of the higher patient dose (represented by size-specific dose estimate (SSDE) delivered to a smaller patient relative to a large patient, for the same scanner output (represented by CTDIvol). Here the patient size is represented by the sum of the anteriopostier (AP) and lateral (LAT) patient dimensions in the center of the scan range. In practice, the scanner output would be lower for the smaller patient and the difference in dose (SSDE) would be minimal.

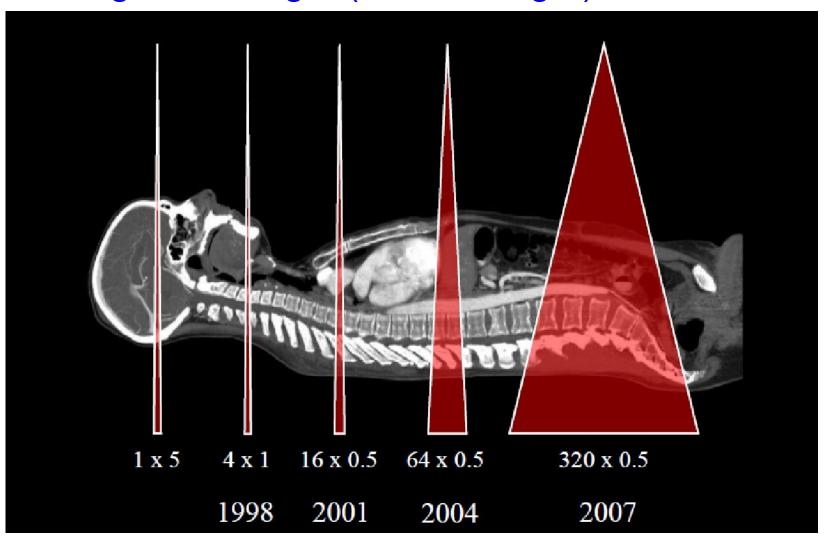


From left: in 545 adult patients undergoing abdominal CT, the scanner output (CTDIvol) increased proportional to the sum of the anterioposterior (AP) and lateral (LAT) dimensions, measured in the center of the scan range.



After conversion to patient dose (SSDE), there is no statistically significant relationship between patient size and dose, demonstrating that the use of higher scanner output values (CTDIvol) in larger patients (to obtain adequate image quality) does not necessarily translate to increased patient dose in larger patients.

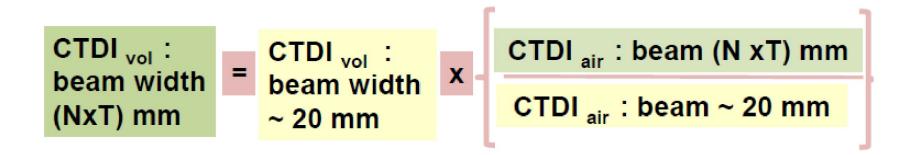
CTDI is not patient dose The integration length (≡ scan length) is not 100 mm

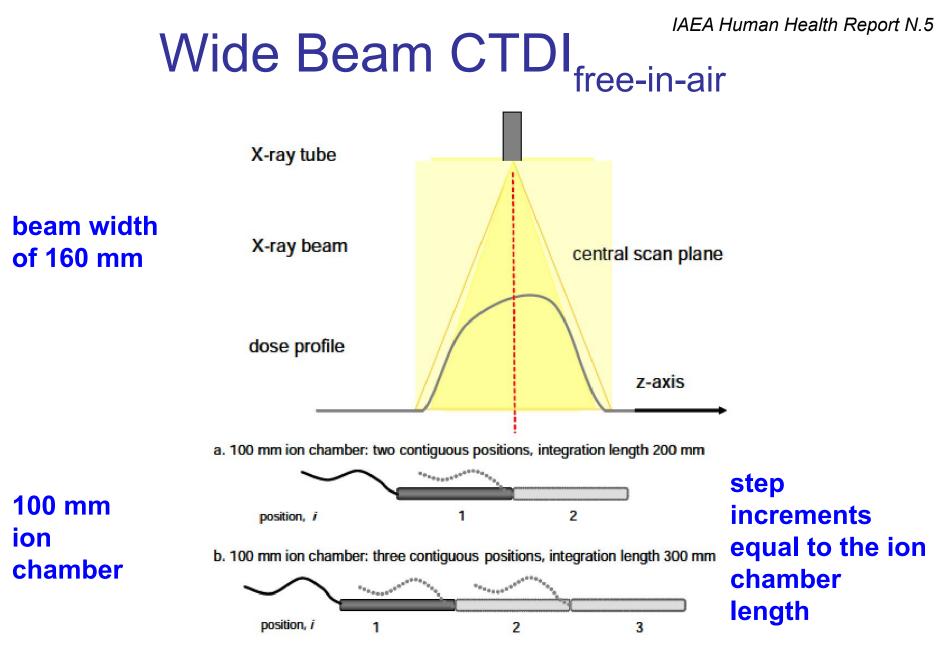


IAEA Human Health Report N.5

IEC: Wide Beam CTDI_{vol}

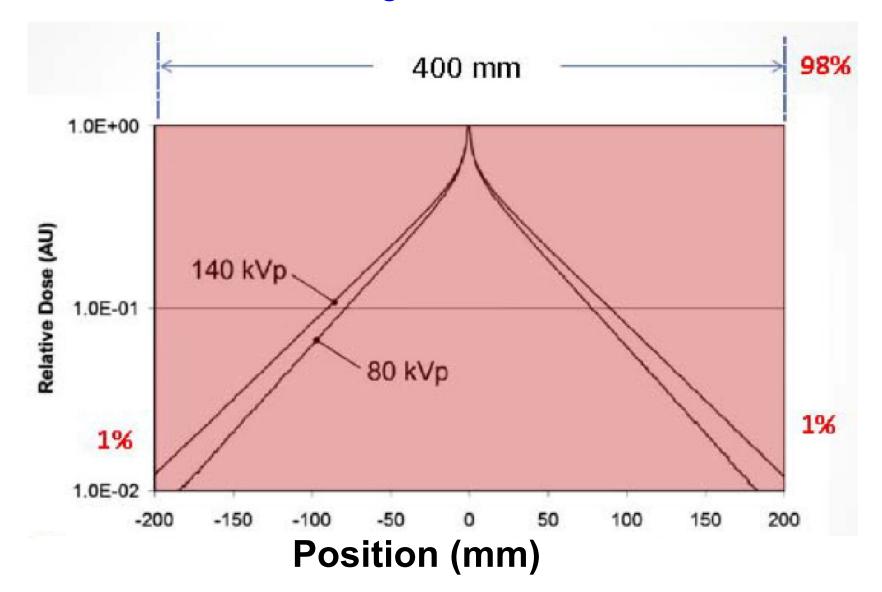
- **CTDIvol** :nominal beam widths (NxT) < 40 mm no change
- CTDIvol :nominal beam widths (NxT) greater than 40 mm
 Measure for an ~ 20 mm beam, correct with CTDI_{air} ratios





The 200 mm integration length is sufficient according to the minimum requirement of IEC, however the 300 mm integration length can also be used

CTDI is not patient dose The scan length is > 100 mm



JM Boone Dose spread function in computed tomography. A Montecarlo Study. Med Phys 36, 2009

AAPM TG111

AAPM REPORT NO. 111



Comprehensive Methodology for the Evaluation of Radiation Dose in X-Ray Computed Tomography

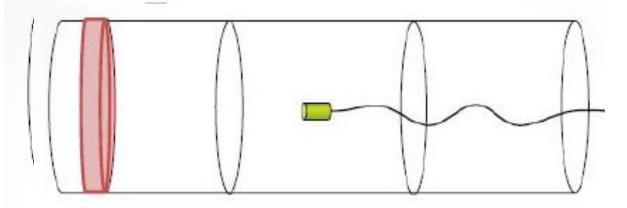
A New Measurement Paradigm Based on a Unified Theory for Axial, Helical, Fan-Beam, and Cone-Beam Scanning With or Without Longitudinal Translation of the Patient Table

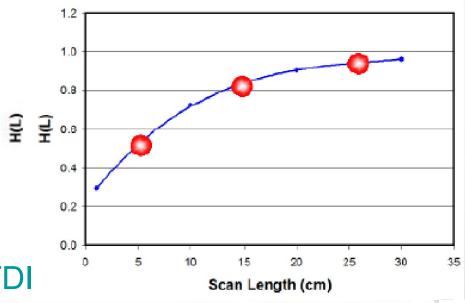


ICRU / AAPM (TG-200) Dosimetry Phantom



AAPM TG111 TG-111 Method





CTDI infinity – equilibrium CTDI

DRLs of CT procedures dedicated for treatment planning in radiation oncology

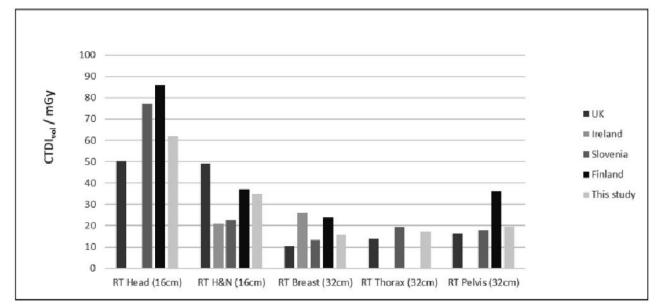
National reference levels of CT procedures dedicated for treatment planning in radiation oncology

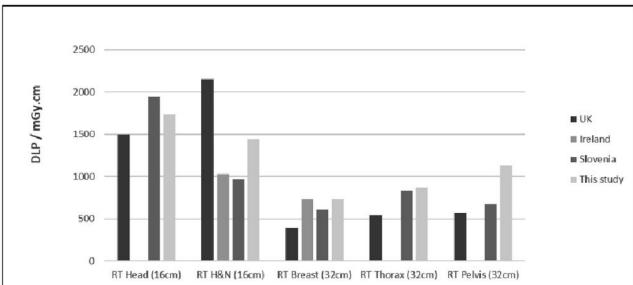
Ana Božanić ^{a, b, *}, Doris Šegota ^a, Dea Dundara Debeljuh ^{a, b, c}, Manda Švabić Kolacio ^a, Đeni Smilović Radojčić ^{a, b}, Katarina Ružić ^d, Mirjana Budanec ^e, Mladen Kasabašić ^f, Darijo Hrepić ^g, Petra Valković Zujić ^{h, i}, Marco Brambilla ^j, Mannudeep K. Kalra ^{k, l}, Slaven Jurković ^{a, b}

| Protocol | Proposed national RPRL | | Achievable level | |
|----------------------|--------------------------|------------|--------------------------|------------|
| | CTDI _{vol} /mGy | DLP/mGy.cm | CTDI _{vol} /mGy | DLP/mGy.cm |
| RT Head (16 cm) | 62 | 1738 | 60 | 1569 |
| RT H&N (16 cm) | 35 | 1444 | 32 | 1422 |
| RT Breast (32 cm) | 16 | 731 | 8 | 361 |
| RT Thorax (32 cm) | 17 | 865 | 9 | 697 |
| RT Pelvis (32 cm) | 20 | 1133 | 17 | 802 |

Proposed national RPRL and achievable national levels for radiation oncology CT acquisitions of the investigated anatomical regions.

DRLs of CT procedures dedicated for treatment planning in radiation oncology





Bozanic et al Phys Med 2022

Can CT protocols used for RT TPS be adjusted to optimize image quality and patient dose?

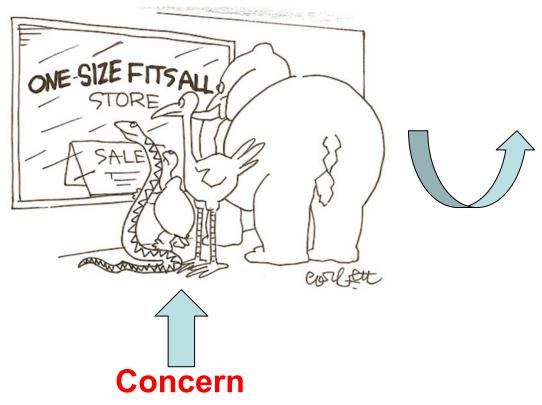
CT images used in RT must serve two key purposes:

- 1. Accurately identify the position of the tumour and OARs
- 2. Provide a map of the ED for the various tissues to be used in the TPS dose calculation

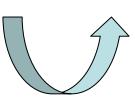
Most radiotherapy centres now have access to dedicated CT scanners. Therefore, the opportunity exists to optimize scan protocol.

On radiotherapy CT scanners a "one size fits all" approach is often taken with minimal variation in scan parameters

Can CT protocols used for RT TPS be adjusted to optimize image quality and patient dose?



Varying scan parameters will change HU values in the images and subsequently introduce inaccuracies to the dosimetric information produced in the TPS.



Disadvantages

Inaccuracies and variability in the outlining process done manually

The use of auto contouring systems might be compromised

Technological developments such as:

- metal artefact reduction
- dual energy imaging
- iterative reconstruction
- TCM
- Automatic KV selection are not used



Problem

Which is the level of HU variation which can be tolerated for different CT imaging techniques, without adversely affecting the dose distribution in the planning process?

Tolerance of HU in guidance documents

| Tissue type | References | RED value | Defined RED or HU tolerance | Corresponding HU ^a |
|-------------|-------------------------------|-----------|-----------------------------|-------------------------------|
| Lung | ESTRO, SGSMP ^{34,35} | 0.2 | ±0.05 (±25%) | ±50 |
| | IPEM ³² | 0.2 | ±0.004 (±2%) | ± 4 |
| | IPEM ³² | 0.4 | ±0.008 (±2%) | ±8 |
| | IAEA ^{23,30,31} | 0.21 | ±0.02 (±10%) or 20 HU | ±20 |
| | AAPM ⁴⁶ | 0.2 | ±50 HU | |
| Soft tissue | ESTRO, SGSMP ^{34,35} | 1.0 | ±0.05 (±5%) | ±50 |
| | IPEM ³² | 1.0 | ±0.01 (±1%) | ±10 |
| | IAEA ^{23,30,31} | 1.06 | ±0.02 (±2%) or 20 HU | ±20 |
| | AAPM ⁴⁶ | 1.0 | ±30 HU | ±30 |
| Bone | ESTRO, SGSMP ^{34,35} | 1.5 | ±0.1 (±7%) | ±170 |
| | IPEM ³⁴ | 1.3 | ±0.03 (±2%) | ±50 |
| | IPEM ³⁴ | 1.8 | ±0.04 (±2%) | ±70 |
| | IAEA ^{23,30,31} | 1.6 | ±0.02 (±1%) or 20 HU | ±34 |
| | AAPM ⁴⁶ | 1.3 | ±50 HU | - |

AAPM, American Association of Physicists in Medicine; ESTRO, European Society for Radiotherapy and Oncology; HU, Hounsfield unit; IAEA, International Atomic Energy Agency; IPEM, Institute of Physics and Engineering in Medicine; RED, relative electron density; SGSMP, Swiss Society for Radiobiology and Medical Physics.

^aHU tolerance calculated using Thomas²⁴ equations.

Scan parameters and level of Hounsfield unit (HU) change in published papers

| CT scan parameter | Impact on HU and scanner manufacturers covered by review | |
|---------------------------|---|--|
| Tube current | No change unless very low current used—GE, Toshiba (Toshiba Medical, Zoetermeer, Netherlands) ^{42,52,53,57} | |
| Kilovoltage | Significant level of HU change—Philips, Toshiba, GE, Siemens (Philips, Amsterdam, Netherlands) ^{2,42,52,53,57} | |
| Acquisition FOV | Depends on CT scanner make/model and which FOV is selected—GE, Toshiba (GE Healthcare, Milwaukee) ^{53,57,58} | |
| | Standard FOVs-no information in articles reviewed | |
| Reconstruction FOV | Extended FOVs—significant change across FOV—Philips, GE ^{59,60} | |
| Slice thickness | Minimal change—Toshiba ⁵⁶ | |
| X-ray tube rotation time | Minimal change—Toshiba ⁵⁶ | |
| Spiral vs sequential | Minimal change—Toshiba ⁵⁶ | |
| Reconstruction algorithms | Depends on CT scanner make/model and which algorithm is selected—Siemens, Toshiba (Siemens Healthcare, Erlangen, Germany) ^{38,56} | |

FOV, field of view.

Summary of HU tolerances to achieve a 1% dose change limit

± 20 HU for soft tissue IAEA± 50 HU for lung and bone AAPM

Note: effects of changes must be considered for all tissue types (air, bone, soft tissue) together when present in the clinical plan.

- 1. a given change of HU or RED will result in a larger change in dose for a greater thickness of tissue than for reduced tissue thickness;
- 2. a single-field treatment plan will deliver a greater dose change for a specific HU change than a multiple field plan;
- 3. the use of lower energy treatment beam results in a higher dose change for a given HU change than the use of higher energy treatment beam;

Advantages of defined tolerance for HU variation during optimization

- When adjusting CT scan protocols, it is helpful to know quickly whether changes to scan protocols are likely to be detrimental to the dosimetric aspects of the planning
- Scan.
- HUs can be easily measured with a phantom on the scanner, thereby allowing early exclusion of inappropriate adjustment to scan parameters.
- Both image quality and HU changes could be assessed with a multipurpose phantom before undertaking a more detailed check to assess the level of dose change in the TPS with an anthropomorphic phantom.
- Use phantoms which approximately match the size and shape of patients when measuring HUs

Conclusions

Scan protocol settings affect image quality. The radiation dose delivered from CT imaging must also be considered and justified.

- The impact of scan parameters in radiotherapy CT is not well detailed in the literature, also considering the number of scanners and the variety of settings within CT protocols
- Publications tend to look at a limited set of scan parameters and only give detailed information on variability when it is considered significant.
- No publications were found which fully assessed the performance of a radiotherapy CT scanner based on variation in both image quality parameters and HU or RED.