

Is there Correlation between Measured and Console $CTDI_{vol}$ in Multislice CT Scanners?

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ABSTRACT

This study investigated the relationship between measured and console volume computed tomography dose index ($CTDI_{vol}$) in multislice CT scanners. The measured $CTDI$ values were determined from kerma length product measured using pencil ionization chamber inserted in the holes of standard CT phantoms with diameters 16 and 32 cm, which respectively mimic adult head and body using the procedures recommended by International Electro-technical Commission. Significantly large deviations were observed between the measured and console $CTDI_{vol}$ values in multislice CT scanners.

Keywords: CT (Computed Tomography); $CTDI$ (Computed Tomography Dose Index); Phantom

INTRODUCTION

Computed Tomography Dose Index ($CTDI$) was introduced for dose optimization and quality assurance in axial CT [1]. In the axial CT, this approach determined, to a good approximation, the average dose for a series of scans. However, the introduction of spiral and multislice CT made the use of this approach in the modern CT designs debatable. While some studies claim that the $CTDI$ approach can also be used in multislice CT, others claim on the contrary. The later argue that as modern designs of multislice CT scanners tend to increase in the scan speed and beam widths, the 100 mm ionization chambers can no longer be used to characterize the entire beam from a single scan profile. The recent development of automatic tube current modulation systems that have been designed to compensate for changes in attenuation will not delineate whether or not the dose is being delivered appropriately over the scan length. In order to ensure confidence in the dose optimization and quality assurance the $CTDI$ approach requires continuous modification to accommodate the technological advancements. This is possible because even after the introduction of spiral CT the $CTDI$ was modified to take into account the effect of pitch [2-4].

The Computed Tomography Dose Index ($CTDI$) as the traditional quantity for dose estimation in CT examinations expressed in mGy, is calculated as described elsewhere [5]:

$$CTDI = \frac{1}{nT} \int_{-50}^{+50} K(z) dz \quad \dots\dots\dots (1)$$

where nT is the total beam collimation (mm) with n the number of active detector rows, T is the nominal slice collimation (mm) and $K(z)$ is the Kerma-Length Product (KLP), which is defined as the product of kerma and thickness of tomographic sections scanned for one gantry rotation. KLP is measured using a 100 mm-pencil ion chamber positioned free in the air or inside the centre and periphery holes of the standard CT phantoms. The phantoms are homogeneous cylinders of diameters 16 cm for head and 32 cm for body made of Polymethylmethacrylate (PMMA). Since the periphery $CTDI$ values are approximately twice those at the centre, the weighted $CTDI$ value ($CTDI_w$) is of interest and is calculated as described elsewhere [5]:

$$CTDI_w = \frac{1}{3}CTDI_{centre} + \frac{2}{3}CTDI_{periphery} \quad \dots\dots\dots (2)$$

$CTDI_w$ gives estimate for the average dose delivered in a number of tomography sections exposed for a single axial scan mode. In order to represent radiation dose for spiral mode it is essential to take into account gaps or overlaps between the X-ray beams during consecutive rotations of the X-ray source. To do this, volume $CTDI_w$ denoted as $CTDI_{vol}$ is calculated as described elsewhere [5]:

$$CTDI_{vol} = CTDI_w \times \frac{1}{pitch} \quad \dots\dots\dots (3)$$

This quantity represents the average absorbed dose over the x, y and z directions of the scanned region within 100 mm length of the ion chamber. However, the $CTDI_{vol}$ remains unchanged whether the scan coverage is equal or greater than 100 mm [5].

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Received: 03-Jun-2022, Manuscript No. JPCB-22-17744; **Editor assigned:** 06-Jun-2022, PreQC No. JPCB-22-17744 (PQ); **Reviewed:** 20-Jun-2022, QC No. JPCB-22-17744; **Revised:** 27-Jun-2022, Manuscript No. JPCB-22-17744 (R); **Published:** 04-Jul-2022, DOI: 10.35248/2161-0398.22.12.337.

Citation: Mhagama AN (2022) Is there Correlation between Measured and Console $CTDI_{vol}$ in Multislice CT Scanners? J Phys Chem Biophys. 12:337.

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To comply with the International Electro-technical Commission (IEC) recommendations all CT scanners display CTDI quantities such as weighted CTDI and volume CTDI for dose optimization and quality assurance [6,7]. CTDI was introduced as a dose descriptor in traditional axial CT scans. Some technological developments including spiral and multislice CT have opened a debate on whether the CTDI approach is still appropriate in the modern designs. In order to establish the suitability of the traditional CTDI approach in spiral multislice CT, this work conducted a validation study to compare the measured and console CTDI_{vol} in spiral multislice CT scanners.

MATERIALS AND METHODS

The study was conducted using six spiral multislice CT scanners acquired by six different hospitals. Information about the scanners types, models are summarized in Table 1. For the sake of simplicity in the presentation of data, the CT scanners have been coded.

Table 1: Summary of CT scanner types and models used in this study.

CT	CT type	Type and model of CT scanner	Manufacturer
SE6C	Siemens	Somatom Emotion 6	Siemens (China)
PB64	Phillips	Brilliance 64	Philips (USA)
SE6G	Siemens	Somatom Emotion 6	Siemens HealthCare (German)
SS16	Siemens	Somatom Sensation 16	Siemens HealthCare (German)
SP128	Siemens	Somatom Perspective 128	Siemens Medical (Shanghai China)

Determination of CTDI according to Equation 1 requires Kerma-Length Product (KLP) values, measured free in air or in standard CT phantoms. These measurements were conducted using SE6C, PB64, SS16, SP128 and SE6G CT scanners, based on hospitals' routine protocols for one rotation of the gantry. These measurements were made using a 100 mm-pencil Unfors Xi CT ion chamber (Serial No. 177013) that was connected to Unfors Xi electrometer. The Unfors electrometer had the uncertainty of 5%, calibration beam quality was RQT, maximum relative deviation for kVp of 3% and energy dependence <5%. The calibration factors were traceable to the international measurement systems [8].

To conduct KLP measurements free in air, the ion chamber was mounted on its stand and positioned free in the air in the scan plane in such a way that the middle of the active region of the chamber, indicated by the shaded part in Figure 1a, was at the centre of the gantry as shown in Figure 1b. The topogram or surview of the ion chamber was then acquired, the scan range (in the middle of the active region of the chamber) selected, the routine head protocol selected, and the scan initiated. The scan parameters, console CTDI_{vol} and KLP values were recorded in the CTDI data sheets such as shown in Figure 1. Additional free-in-air KLP measurements for routine chest, abdomen and pelvis protocols were made using similar procedures but with the automatic tube current modulation system deactivated (Table 2).

Measurements of KLP in the head phantom, shown in Figure 2a right, was made using routine head protocol with the ion chamber inserted in five holes (one at a time) labeled center (C), North (N), East (E), South (S) and West (W) shown in Figure 2b. In each turn of the hole measurements, the four remaining holes had their inserts in place.

With the ion chamber inserted into one of the holes as shown in Figure 3a, the head phantom was placed in the scan plane as shown in Figure 3b. The topogram or surview of the phantom was then acquired, the scan range (at the middle of the phantom) selected, routine head protocol also selected and the scan initiated.

The scan parameters used and the KLP values obtained were recorded in the CTDI data sheets such as shown in Table 2. The ion chamber was then replaced into another hole to acquire additional KLP values and the same procedures were repeated until measurements in all five holes were made. KLP measurements in phantom under routine chest protocol were made using the body phantom shown in Figure 2a. The same procedures, including the positioning of the phantom and ion chamber, the acquisition of the topogram or surview and the selection of routine protocol were made as described during KLP measurements in the head phantom. However, in the chest measurements, after the acquisition of topogram or surview and the selection of the scan range, the AEC system was deactivated then the scan initiated. The same procedures were repeated to obtain KLP values for routine abdomen and pelvic protocols.

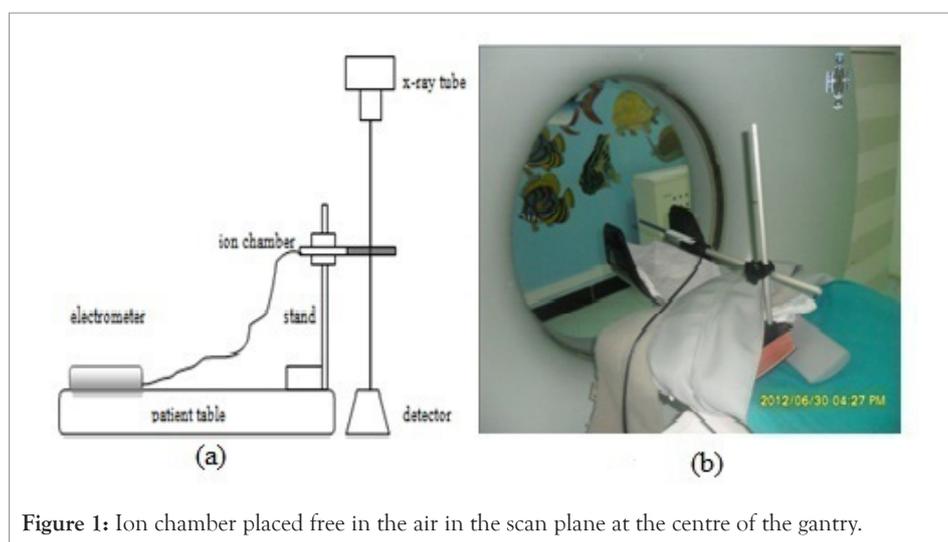


Figure 1: Ion chamber placed free in the air in the scan plane at the centre of the gantry.

Table 2: CTDI data sheet for routine head protocol using SS16.

TMJ routine protocol	Chamber position	Tube potential (kV)	Tube current (mAs)	Rotation time (s)	Beam collimation (mm)	Slice thickness (mm)	Table feed (mm)	Console CTDI _{vol} (mGy)	KLP (mGy.cm)
Head	Free in air	120	250	1	12 × 0.75	4.5	9	60.48	45.84
								60.48	45.89
	C	120	250	1	12 × 0.75	4.5	9	60.48	42.45
								60.48	42.33
	N	120	250	1	12 × 0.75	4.5	9	60.48	49.19
								60.48	49.04
	W	120	250	1	12 × 0.75	4.5	9	60.48	46.96
								60.48	47.11
	S	120	250	1	12 × 0.75	4.5	9	60.48	44.71
								60.48	44.36
	E	120	250	1	12 × 0.75	4.5	9	60.48	48.06
								60.48	47.81

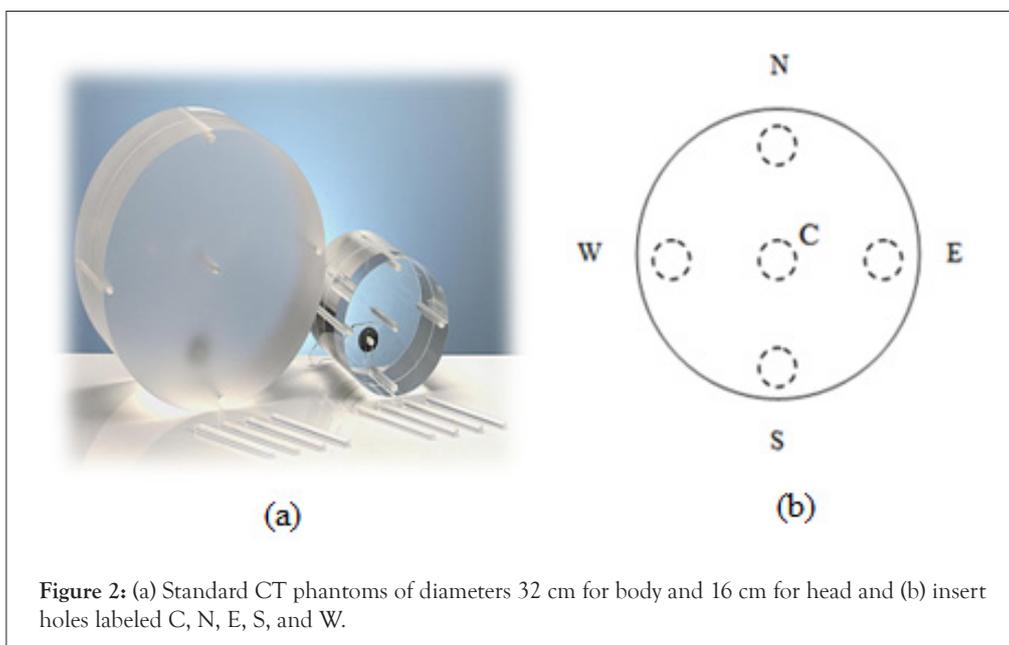


Figure 2: (a) Standard CT phantoms of diameters 32 cm for body and 16 cm for head and (b) insert holes labeled C, N, E, S, and W.

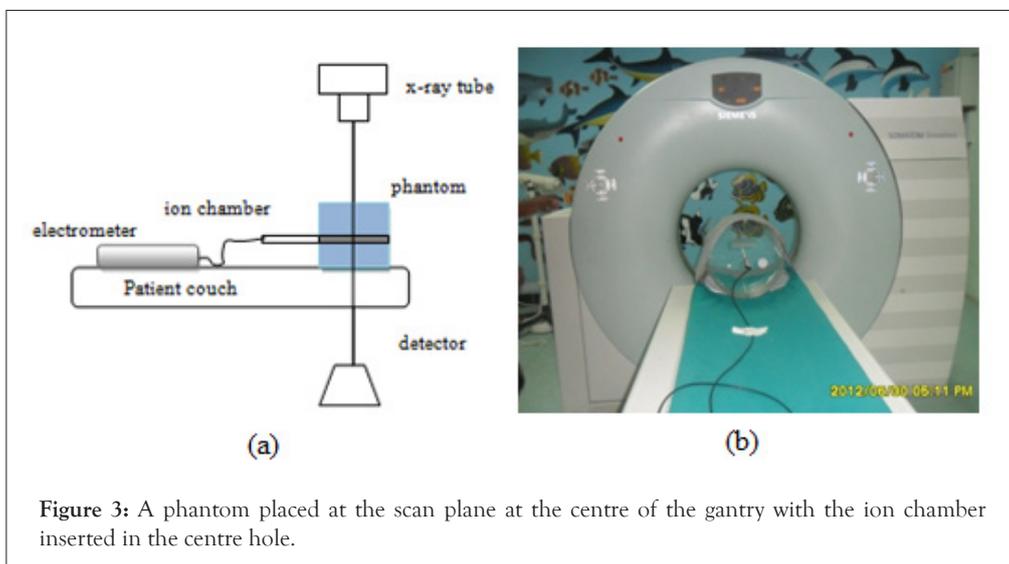


Figure 3: A phantom placed at the scan plane at the centre of the gantry with the ion chamber inserted in the centre hole.

RESULTS AND DISCUSSION

The mean scan parameters used and KLP values obtained during different measurements were extracted from the CTDI data sheets and summarized in Table 3.

The air, centre and periphery CTDI values presented in Table 4 were obtained according to Equation (1) using KLP and beam collimation values extracted from Table 3 as inputs to this equation. Furthermore, the $CTDI_w$ and $CTDI_{vol}$ values also presented in this table were obtained using Equations (2) and (3) respectively. The values of measured and console displayed $CTDI_{vol}$ are also presented in this table. It is evident from this table that for most hospitals and protocols, the periphery CTDI values were as expected about twice the centre CTDI values. The observed deviations in the head protocols could be attributed to scatter radiation that contributed to number photons detected by the ion chamber when inserted in the centre hole of the head phantom.

It was also observed that despite using the same model of CT scanner, SE6C produced higher CTDI values than those obtained at SE6G for all routine protocols. This was attributed to fact that SE6C used higher effective tube loads of 350 mAs and 120 mAs for head and abdomen/pelvis protocols respectively as compared to 267, 100,105 mAs used at SE6G for head, chest/abdomen, and pelvic protocols, respectively. However, for the chest protocol, higher CTDI value observed at SE6C could be attributed to the use of spiral mode with thicker slices (5.0 mm) and wider beam collimation (11.1 mm) as compared to sequential mode used in SE6G with the slice thickness of 1.0 mm and beam collimation of 1.0 mm. Large variations of $CTDI_w$ values were observed among the hospitals. The $CTDI_w$ values for head protocol were the highest at SP128 and the lowest at SS16. The highest head $CTDI_w$ values at SP128 could be attributed to the use of high tube potential (130 kV), long rotation time (2 s), and large slice thickness (10 mm). The

lowest values obtained at SS16 could be explained by the use of relatively lower tube potential (120 kV), shorter rotation time (1.0 s), and smaller slice thickness (4.5 mm).

The data presented in Table 4 for measured and console $CTDI_{vol}$ values for every CT scanners were compared and summarized for routine protocols in Figures 4-8. Generally, these figures present large deviation from the measured and console $CTDI_{vol}$. It is evident from Figures 4a, 5a, and 7a that there are large deviations between the measured and console $CTDI_{vol}$. Non-origin linear relationship observed in Figures 4b, 5b and 7b indicates that there is no correlation between the quantities and therefore the measured and console $CTDI_{vol}$ do not match.

Unlike in SE6G, PB64, and SS16, Figures 6 and 8 show marginal comparison and correlation between the measured and console $CTDI_{vol}$, although it is still not clear what causes these deviations.

The $CTDI_w$ values can be used to assess the relative risks associated with radiation dose by using the dose reference levels presented in the third quartiles and mean values. The third dose quartiles for different protocols set by the European Commission's Radiation Protection Action represent the bounds of potentially unacceptable practice [9]. The third dose quartiles for different protocols and hospitals in Tanzania and the European Commission have been compared in Figure 9.

In general, the $CTDI_w$ values obtained in this study were comparable or less than the corresponding values obtained from the European Commission. Thus, based on the dose reference levels, the risks associated with the radiation dose in CT examinations in Tanzania is tolerable. However, higher $CTDI_w$ value for the head protocol observed in SP128 could be attributed to the use of the large slice thickest of 10 mm as presented in Table 3.

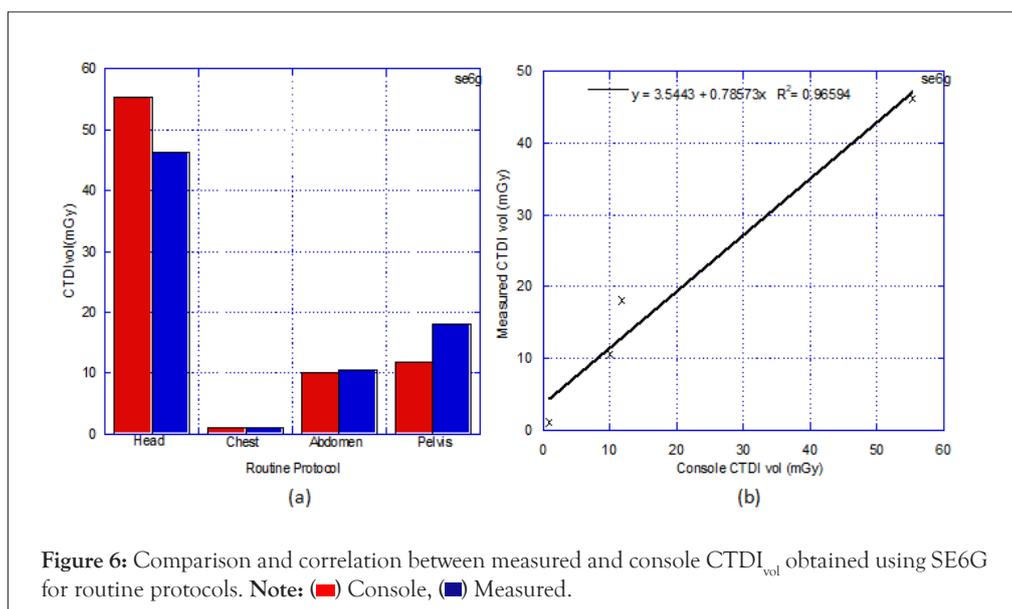
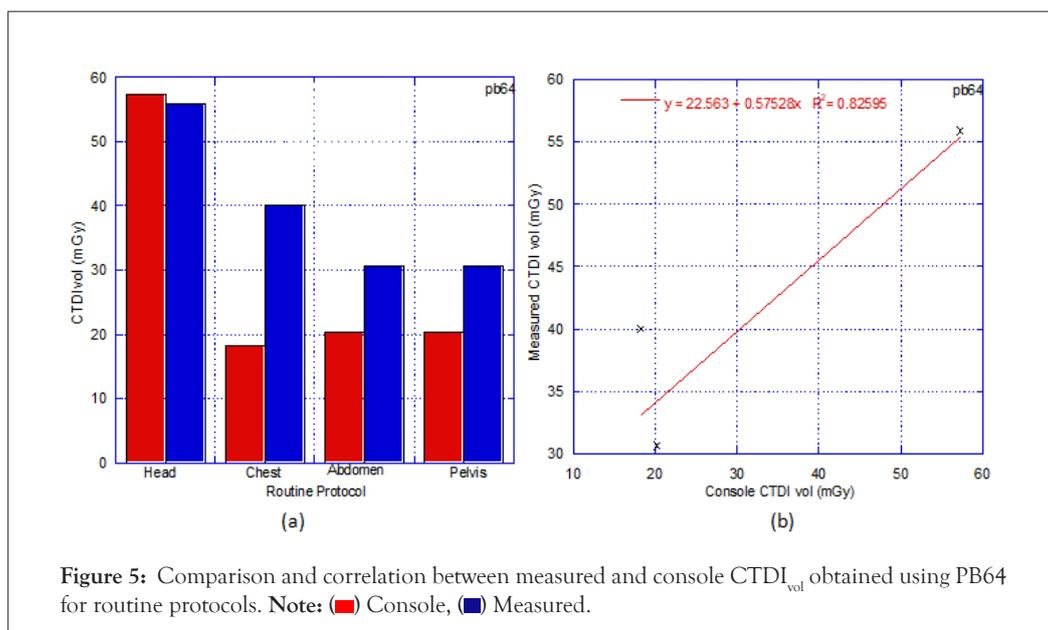
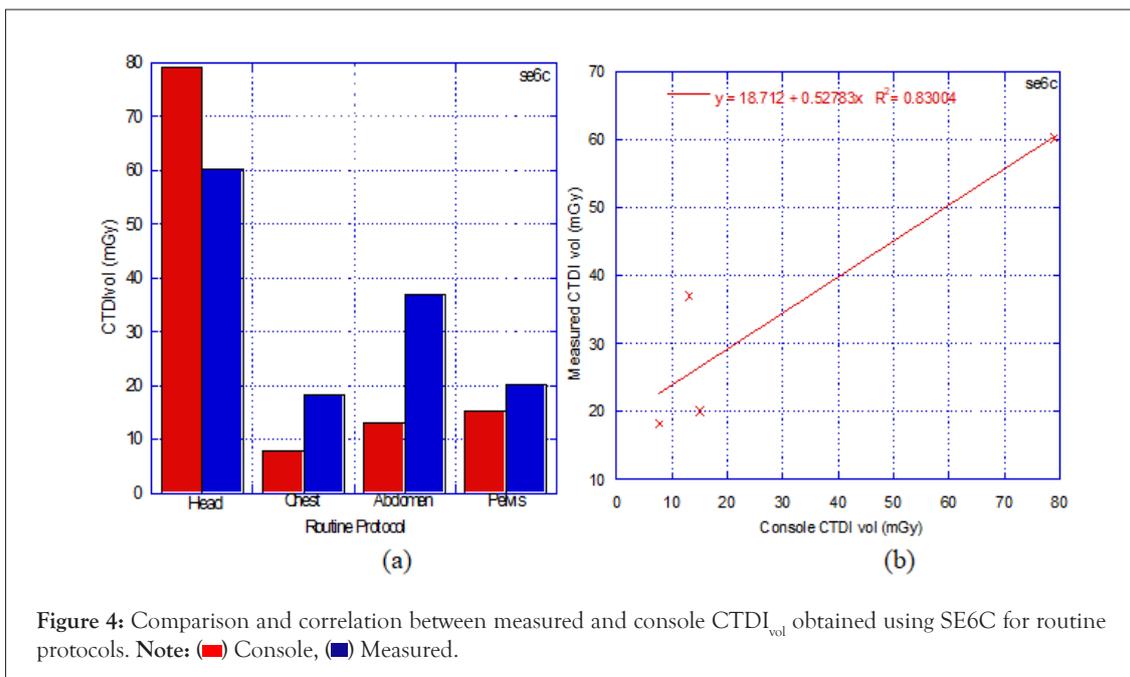
Table 3: Scan parameters used for routine protocols at different hospitals during CTDI determination.

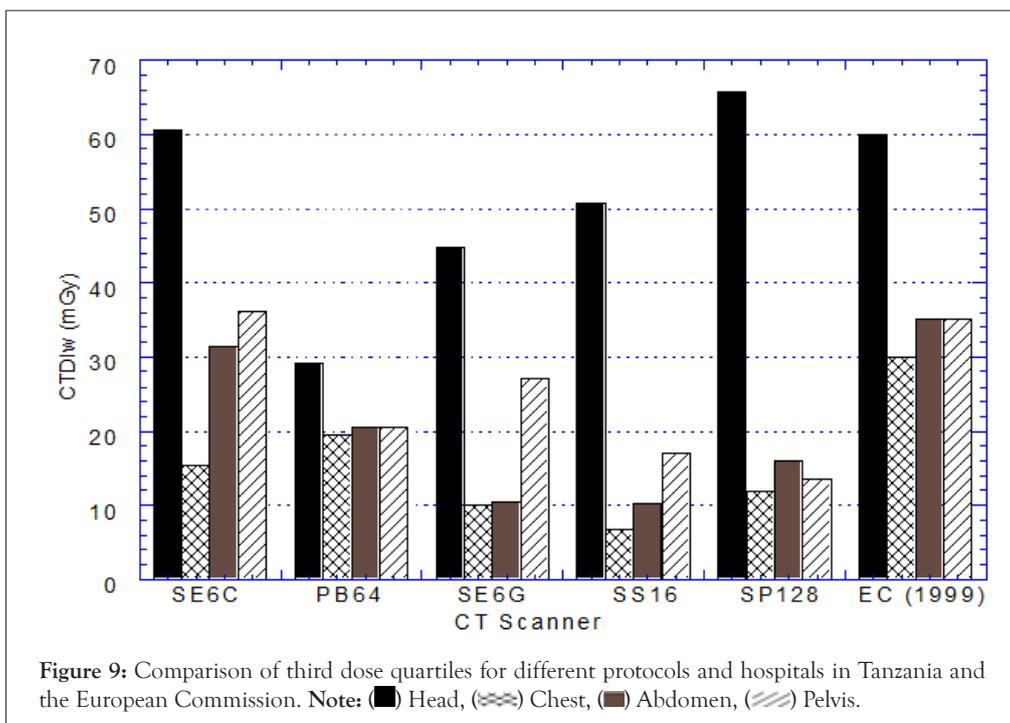
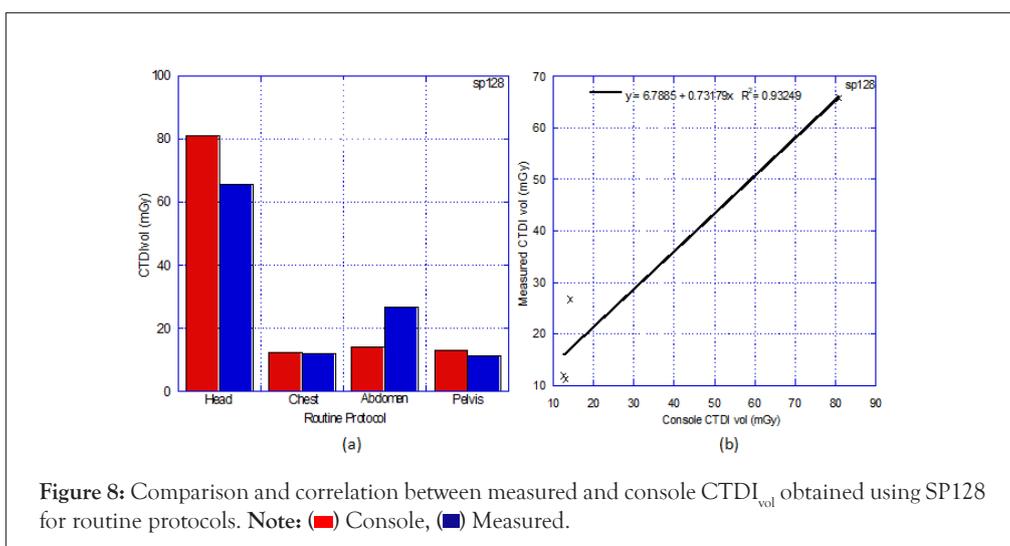
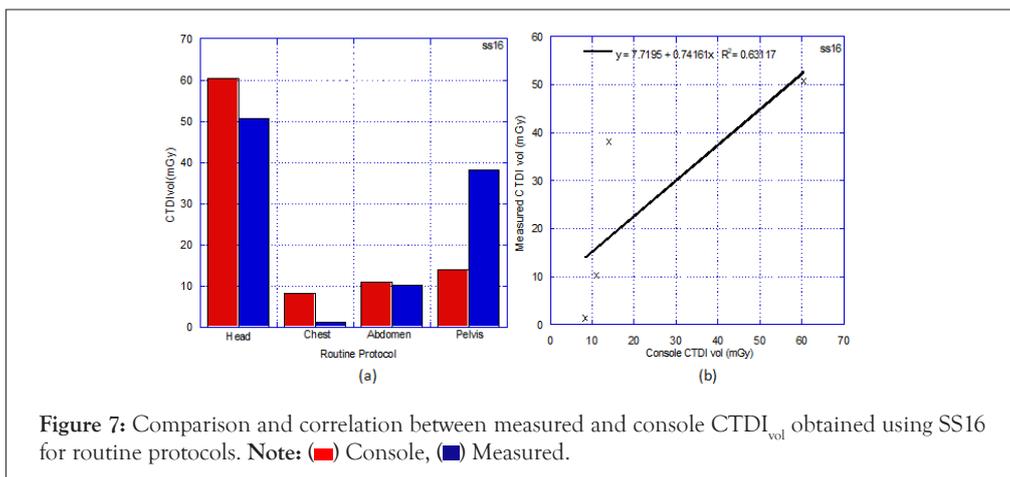
Protocol	CT	Console Readings							Kerma-length product (mGy.cm)			
		Tube potential (kV)	Effective mAs or mAs/slice	Rotation time (seconds)	Slice thickness (mm)	Beam collimation (mm)	Table feed (mm)	Pitch factor	Console $CTDI_{vol}$ (mGy)	Air	Centre	Periphery
Head	SE6C	130	350	1.5	4.9	16.3	16.3	-	79.1	168.5	54.3	120.6
	PB64	120	350	0.8	2	40	-	0.5	57.3	234.4	74.4	138
	SE6G	130	267	1.5	5	10.3	10	-	55.4	9.3	45.3	46.6
	SS16	120	267	1	4.5	9	9	-	60.48	45.8	42.4	47.3
	SP128	130	270	2	10	7.2	7.2	-	80.9	74.4	43.8	49.1
Chest	SE6C	130	70	0.6	5	11.1	-	0.9	7.7	63.5	9.2	21.1
	PB64	120	300	0.8	0.9	40	-	0.5	18.3	86.4	33.2	40
	SE6G	130	100	1	1	1	10	-	1	2.8	0.6	1.2
	SS16	120	100	1	1	2	10	-	8.4	2.7	0.8	1.7
	SP128	130	100	1	1	2	2	-	12.6	7.4	1.4	2.9

Abdomen	SE6C	130	120	0.6	5	12	-	0.9	13.1	122	23.3	45
	PB64	120	250	0.8	0.9	40	-	0.7	20.3	99.2	62.8	91.6
	SE6G	130	100	0.8	5	10	10	-	10	12.7	5.3	13.2
	SS16	120	175	0.8	5	10	10	-	11	20.4	5.9	12.5
	SP128	130	130	0.6	5	38.4	-	0.6	14.3	179.7	34.2	74.9
Pelvis	SE6C	130	120	1	3	6	-	1.8	15.2	30	10.4	27.3
	PB64	120	250	0.8	0.9	40	-	0.7	20.3	99.2	62.8	91.6
	SE6G	130	105	1	5	12	-	1.5	11.9	58.5	19.1	39
	SS16	120	200	0.5	5	24	-	0.5	14	84.2	23.8	49.9
	SP128	130	120	1	5	38.4	-	1.2	13.2	242.3	35.3	59.9

Table 4: Measured and console CTDI values for head and body using routine protocols.

Protocol	Hospital	CTDI _{air}	CTDI _{centre}	CTDI _{periphery}	CTDI _w	Measured CTDI _{vol} (mGy)	Console CTDI _{vol} (mGy)
Head	SE6C	103.4	33.3	74	60.5	60.3	79.1
	PB64	58.6	18.6	34.5	29.2	55.9	57.3
	SE6G	9	44	45.2	44.8	46.2	55.4
	SS16	50.9	47.1	52.5	50.7	50.7	60.48
	SP128	103.4	60.9	68.2	65.8	65.8	80.9
Chest	SE6C	57.2	8.3	19	15.4	18.2	7.7
	PB64	21.6	8.3	10	19.4	40	18.3
	SE6G	27.6	5.5	12.3	10	1	1
	SS16	13.4	3.9	8.3	6.8	1.4	8.4
	SP128	37.2	6.9	14.3	11.9	11.9	12.6
Abdomen	SE6C	101.7	19.4	37.5	31.4	37	13.1
	PB64	24.8	15.7	22.9	20.5	30.6	20.3
	SE6G	12.7	5.3	13.2	10.5	10.5	10
	SS16	20.4	5.9	12.5	10.3	10.3	11
	SP128	46.8	8.9	19.5	16	26.7	14.3
Pelvis	SE6C	50	17.4	45.5	36.1	20.1	15.2
	PB64	24.8	15.7	22.9	20.5	30.6	20.3
	SE6G	48.8	15.9	32.5	27	18	11.9
	SS16	35.1	9.9	20.8	17.1	38.1	14
	SP128	63.1	9.2	15.6	13.5	11.3	13.2





In order to make similar comparison, the $CTDI_w$ values for each protocol, were averaged to obtain the mean and third quartile $CTDI_w$ values for this study. These values were compared in Table 5 with the corresponding values obtained from other

studies. It is evident from this table that, the mean and third quartile $CTDI_w$ values obtained in this study were comparable to those obtained from other studies.

Table 5: Mean and third quartile CTDI_w values (mGy) for different protocols obtained in this study compared with different studies.

Routine protocol	This study		[10]		[11]		[12]		[9]	
	Mean	Third quart	Mean	Third quart	Mean	Third quart	Mean	Third quart	Mean	Third quart
Head	50.2	60.5	42.9	53.8	55	62	22.5	-	50	60
Chest	12.7	15.4	16.5	18.4	20	21	12.6	-	20.3	30
Abdomen	17.7	20.5	17.1	18.4	22	23	11.3	-	25.6	35
Pelvis	22.8	27	18.5	22.6	22	23	11.3	-	26.4	35

In order to facilitate CT performance tests, the International Electro-technical Commission recommended that all CT scanners should display, on their consoles, the radiation dose quantities in terms of CTDI_w or CTDI_{vol} after patient examination [7-12]. In view of this requirement, this study was interested to compare the calculated and the console CTDI_{vol} values presented in Tables 3 and 4, respectively. Large deviations were observed between the calculated and console CTDI_{vol} values. These deviations could be attributed to different techniques used to obtain the calculated and console CTDI_{vol} values. This is because, during KLP measurements, the CT scanners displayed the fixed CTDI_{vol} values on their consoles for every KLP measurements made, despite the changes in the ion chamber positioning in the insert holes of the standard CT phantoms. As presented in Table 2, the console CTDI_{vol} value was fixed at 60.48 mGy in all ion chamber positions during KLP measurements in the head phantom. For these measurements, this study calculated the CTDI_{vol} value using Equation (3) to obtain 50.7 mGy as presented in Table 4. The unclear definition of CTDI_w and CTDI_{vol} by the CT manufacturers, could explain why there is limited standardization between the console and the calculated CTDI_{vol}. It was difficult for this study, therefore, to establish the correlation between the console and the calculated CTDI_{vol}.

CONCLUSION

The assessment of CTDI in computed examinations in different CT scanners showed that, on average, multislice CT scanners delivered comparable radiation doses are within the European Commission's diagnostic reference levels and therefore tolerable. Despite the lower levels of radiation dose, there is significant deviation between the calculated and console CTDI in all spiral multislice CT scanners studied. This explains why the CTDI technique for quality assurance and dose optimization is no longer appropriate for modern spiral multislice CT

scanners with high rotation speeds and wide beams as discussed elsewhere. The CTDI approach for dose optimization and quality assurance should be continuously modified to fit with the CT technology developments.

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