



Sudan University of Science & Technology
College of Graduate Studies



ESTIMATION OF RADATION DOSE FOR PATIENT CHEST DURING COMPUTED TOMOGRAPHY

تقدير الجرعة الإشعاعية للمرضى أثناء التصوير التصوير المقطعي للصدر

A thesis submitted for partial fulfillment for the requirement of MSc degree in
Medical Physics

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2020

الآية

(أَمَّنْهُ قَانِشَ آتَاءَ اللَّيْلِ سَاجِدًا وَقَانِمًا يَحْذَرُ الْإِخْرَةَ وَيَرْجُو
رَحْمَةَ رَبِّهِ ۖ قَدْ هَذَا يَسْتَوِي الْأَنِينَ يَعْلمُونَ وَالنَّيْنَ لَهُ يَعْلمُونَ ۖ

إِنَّمَا يَتَذَكَّرُ أُولُو الْأَلْبَابِ) [الزمر الآية (9)]

Acknowledge

First of all, I thank Allah the Almighty for helping me complete this research. I thank **Dr. Hussein Ahmed Hassan** my supervisor, for his help and guidance.

Finally I would like to thank everybody who helped me preparing and finish this study.

Dedication

DEDICATE THIS RESEARCH TO MY
PARENT MAY GOD HAVE MERCY AND
THANKS, MY SONS , MY BROTHERS AND
MY SISTERS , MY FRIENDS AND THE
SUPERVISOR WHO STAND ME AND
SUPPORT ME IN MY EDUCATION
CAREER

Abstract

The main of study to estimate the radiation dose for the patients chest (CT) examination undergoing (CT) scanning .

This study in Khartoum – State Modern Medical Center the period December2019. The data include 50 patients male and female.

Use machine CT GE OPTIMA 60slices.

Method include to measure the dose radiation which controlmany parameters (Kvp, MAs,CTDL_{vol}, DLP)this parameter was calculated using of machine after complication exam done.

Use estimated (DLP) and effective does (E).

Analysis of data by (spss) In additional should result analysis the mean effective does for achest (11.29) (mSv) the mean (DLp) 774.76 mGY/cm. and the mean (CTDLvol) (4.620mGY) .

The Maximum Dose(29mSv) and the Minimum(2mSv).

(CT) accounts for the majority of radiation exposure related to medical imaging medical professionals should have a working knowledge of the benefits and risk of medical radiation and an understand of strategies for reducing (CT) radiation .

المستخلص

تهدف الدراسة إلى تقدير الجرعة الإشعاعية لصدر المرضى اثناء التصوير المقطعي CT(scan)

لعينة مكونة من 50 مريض خضعوا لامتحانات التصوير المقطعي المحوسب بالصدر في المركز الطبي الحديث بولاية الخرطوم.

وتم استخدام CTDL(vol) وطول الجرعة (DLP) والجرعة الفعالة (E).

تم تحليل البيانات عن طريق برنامج التحليل الإحصائي المعروف (SPSS) لمعرفة المتوسط الحسابي والانحراف المعياري.

أظهرت نتائج التحليل متوسط الجرعة الفعالة ومتوسط الجرعة $E \text{ mSv}$ (11.29) ومتوسط $CTDL(vol)$ (4.620 mGY) ومتوسط DLP (774.76 mGY/cm).

اعلى جرعة (29 mSv) واقل جرعة (2mSv).

تقدير الجرعة الاشعاعية المعرضة باستخدام CT يجب ان تحدد بواسطة شخص مسؤول له علاقة بالتصوير الطبي وان يكون علي معرفة الفوائد والمخاطر والاشعاع الطبي ودراسة طرق تحليل الجرعة الاشعاعية في CT (scan).

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List of abbreviation

CT	Computer Tomography
ALARA	As Low As Reasonably Achievable
ICRP	International Commission on Radiological Protection
DRLs	Diagnostic Reference Dose Levels
QA	Quality assurance
CAT	Computerized axial Tomography
DAS	Data Acquisition System
MSAD	Multiple Scan Average Dose
MDCT	Multi Detector Computer Tomography
CTDI	Computer Tomography does index
CTDI ₁₀₀	Computer Tomography does index, for a100 mm length pencil ion chamber
CTDI _{vol}	Volumetric Computer Tomography does index
CTDI _w	Weighted Computer Tomography does index
DLP	Dose Length Product
E	Effective Dose
D	Absorbed Dose
FDA	Food and Drug Administration
WR	Radiation Weighting Factor
WT	Tissue Weighting Factor
KERMA	Kinetic Energy Released per unit Mass
Kvp	Kilo Voltage Peak
mAs	Mile Ampere Second
NRPB	National Radiological Protection Board
D _T	Organ Dose
H _T	Equivalent Dose
MeV	Mega electron Volt
mSv	Mile Sievert
EC	European Commission
NRPB	National Radiological Protection Board
ImPACT	Impact CT patient dosimetry calculator
AEC	Automatic Exposure Control

CHAPTER ONE

INTRODUCTION

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INTRODUCTION

1.1 Introduction:

1.1.1 Ionization radiation:

Ionizing radiation is any type of particle or electromagnetic wave that carries enough energy to ionize or remove electrons from an atom. There are two types of electromagnetic waves that can ionize atoms: X-rays and gamma rays, and sometimes they have the same energy. Gamma radiation is produced by interactions within the nucleus, while X-rays are produced outside of the nucleus by electrons. There are officially two types of ionizing radiation that are energetic particles emitted during an interaction within the nucleus.

Ionizing radiation dangerous when atoms in living cells become ionized one of three things usually happen the cell dies, the cell repairs itself, or the cell mutates incorrectly and can become cancerous.

1.1.2 CT Scan:

A CT scan makes use of computer-processed combinations of many X-ray measurements taken from different angles to produce cross-sectional (tomographic) images (virtual "slices") of specific areas of a scanned object, allowing the user to see inside the object without cutting. Other terms include computed axial tomography (CAT scan) and computer aided tomography.

Digital geometry processing is used to further generate a three-dimensional volume of the inside of the object from a large series of two-dimensional radiographic images taken around a single axis of rotation. Medical imaging is the most common application of X-ray CT. Its cross-sectional images are used for diagnostic and therapeutic purposes in various medical disciplines. The rest of this article discusses medical-imaging X-ray CT; industrial applications of X-ray CT are discussed at industrial computed tomography scanning.

The term "computed tomography" (CT) is often used to refer to X-ray CT, because it is the most commonly known form. But, many other types of CT exist, such as positron emission tomography (PET) and single-photon emission computed tomography (SPECT).

CT produces a volume of data that can be manipulated in order to demonstrate various bodily structures based on their ability to absorb the X-ray beam. Although, historically, the images generated were in the axial or transverse plane, perpendicular to the long axis of the body, modern scanners allow this volume of data to be reformatted in various planes or even as volumetric (3D) representations of structures. Although most common in medicine, CT is also used in other fields, such as nondestructive materials testing. Another example is archaeological uses such as imaging the contents of sarcophagi. Individuals responsible for performing CT exams are called radiographers or radiologic technologists

1.2 Problem of Study:

Ratio of disease of chest is very high according the chest upper body muscles its control the ability to perform every day activity so the frequency of the patient chest is high in hospital or clinical. The radiation of chest CT is very sensitive organ like heart, lung.

Computed tomography (CT) chest can involve relating in high dose to patient. The dose can exceed levels known with to increase the probability cancer.

There for the radiation dose measurement during the chest CT test is consider important point in order to mach accepted level of received dose.

1.3 objective of Study:

1.3.1: general objective:

The objective of study was to determine of effective dose by use CT-Scan in medical diagnosis and optimized dose Chest CT.

1.3.2: Specific Objective:

- Estimated Effective dose associated with low-dose Chest CT.
- Evaluate the suitable of radiation dose for adult patient in Chest CT scanning.
- Quantify the patient dose in CT examination for Chest.

1.4: Significant of Study:

The important of this study the radiation dose for chest CT and compare it is dose value with standard international acceptable level of

Chest CT dose, also explain radiation risk during in CT for Chest. And stimulate the effective dose for the patient.

1.5: Over view this Study:

This study was consist of five chapters, chapter one will be an introduction introduce briefly this thesis and it was contain, general introduction about Radiation ionization Risk and CT-scan dose measurement and radiation risk, problem of the study, general and specific objective, significant of the study in addition to the overview of the study. Chapters two was the literature review which contain the general theoretical background of CT-scan and study about evaluation of dose and calculate of during computed tomography scan for brain also (effective dose and risk of radiation). Chapter three will describe the methodology (material, methods) was used in this study. Chapter four was including results of presentation of final finding of study. Chapter Five was including discussion, conclusion and recommendation for future scope in addition to references and appendices.

CHAPTER TWO

THEORETICAL BACKGROUND

CHAPTER TWO

THEORETICAL BACKGROUND

2.1 Introduction

Since the introduction of helical computed tomography (CT) in the early 1990s, the technology and capabilities of CT scanners have changed tremendously (helical and spiral CT are equivalent technologies; for consistency, the term “helical” will be used throughout). The introduction of dual-slice systems in 1994 and multislice systems in 1998 (four detector arrays along the z-axis) has further accelerated the implementation of many new clinical applications. The number of slices, or data channels, acquired per axial rotation has increased, with 16- and 64-slice systems now available (as well as models having 2, 6, 8, 10, 32, and 40 slices). Soon even larger detector arrays and axial coverage per rotation (>4 cm) will be commercially available, with results from a 256-slice scanner having already been published⁴. These tremendous strides in technology have resulted in many changes in the clinical use of CT. These include, but are not limited to, increased use of multiphase exams, vascular and cardiac exams, perfusion imaging, and screening exams (primarily the heart, chest, and colon, but also self-referred “wholebody” screening exams). Each of these applications prompts the need for discussion of radiation risk versus medical benefit. In addition, the public press in the United States, following publication by the *American Journal of Roentgenology* of two articles on risks to pediatric patients from CT, has begun to scrutinize radiation dose levels from all CT examinations. Subsequent reports in the popular media have increased the concern of patients and parents of pediatric patients undergoing medically appropriate CT examinations.

the importance of radiation dose from x-ray CT has been underscored recently by the attention given in the scientific literature to issues of dose and the associated risk. The dose levels imparted in CT exceed those from conventional radiography and fluoroscopy and the use of CT continues to grow, often by 10% to 15% per year. According to 2006 data, approximately 62 million CT examinations were performed in hospitals and outpatient imaging facilities in the United States¹⁶. Thus, CT will continue to contribute a significant portion of the total collective dose delivered to the public from medical procedures involving ionizing radiation.

The rapid evolution of CT technology and the resultant explosion in new clinical applications, including cardiac CT, combined with the significance of CT dose levels, have created a compelling need to teach, understand, and use detailed information regarding CT dose.

All of these factors have created a need for the AAPM to provide professional, expert guidance regarding matters related to CT dose. Fundamental definitions of CT dose parameters require review and, perhaps reinterpretation, as CT technology evolves: some parameter definitions are not being used consistently, some are out of date, and some are more relevant than others with respect to patient risk or newer scanner designs. Hence, this task group, consisting of experts in the field of x-ray CT, was formed to address the following:

1. Provide guidance from the AAPM to the International Electrotechnical Commission (IEC) and U.S. Food and Drug Administration (FDA) and facilitate international consensus regarding:

- a. CT dose parameter definitions.
- b. CT dose measurements.
- c. Appropriate use of CT dose and risk parameters

2. Educate AAPM members regarding:

- a. CT dose definitions and measurements.
- b. European Guidelines on Quality Criteria for Computed Tomography, and other relevant international publications.
- c. Reasonable dose levels for routine CT examinations.

2.2 Overview of Multiple-Detector-Row Ct (Mdct) Technology:

The “multiple-detector-row” nature of MDCT scanners refers to the use of multiple detector arrays (rows) along the z -direction (perpendicular to the axial CT plane). Currently available MDCT scanners utilize third-generation CT geometry in which an arc of detectors and the x-ray tube(s) rotate together. All MDCT scanners use a slip-ring gantry, allowing helical acquisition at rotation speeds as fast as 0.33 seconds for a full rotation of the x-ray tube about the isocenter.

These scanners offer tremendous flexibility because of their advances not just in detector technology, but also in data acquisition

systems (DAS), x-ray tube design and other subsystems. One illustration of this is that while MDCT scanners have multiple rows of detectors, the data collected from multiple rows can be combined as though they were collected from one detector. To describe this behavior, the term “channel” has been used, where a channel is the smallest unit in the z -direction from which data are independently collected. Therefore, if data from multiple detector rows are collected in such a way that those individual rows are combined (and the ability

to determine from which row the data was originally collected is lost), then these rows form a channel.

Currently, commercial MDCT systems are capable of acquiring up to 64 channels of data (along the z -direction) simultaneously. Other values for N_{max} , the maximum number of independent channels acquired along the z -axis, for current commercially available MDCT systems include: 2, 4, 6, 8, 10, 16, 32, and 40. For axial data acquisitions, each channel collects sufficient data to create one “slice” or image, so as many as N_{max} independent images along the z -axis could theoretically be reconstructed. For narrow slice widths, however, cone-beam geometrical considerations may limit the number of allowed images per rotation to less than N_{max} . For example, one manufacturer’s 16-slice scanner allows sixteen 1.5-mm channels for helical acquisitions, but only twelve 1.5-mm channels for sequential acquisitions because of cone-beam considerations. Alternatively, the data from multiple channels can be added together to create fewer than N_{max} images per gantry rotation, each image having a relatively wider width. Compared to singledetector- row CT (SDCT), MDCT systems allow faster acquisitions of a volume of data with less heat load on the x-ray tube (both by a factor of up to N_{max}). Sixty-four-channel systems appeared on the market in the middle of 2004, sixteen-channel scanner models were announced at the end of 2001, and four-channel systems were introduced in late 1998. A 256-channel system is anticipated in 2008.

Figure 1 illustrates the difference between an SDCT scanner and an MDCT scanner from the same vendor. The fundamental difference is the use of multiple detector-rows along the z direction. In SDCT, the slice width is primarily determined by the pre-patient x-ray collimation (post-patient collimation was used on some SDCT scanners models). In MDCT, the z -extent of the data acquisition is determined by the pre-patient x-ray

beam collimation, but the slice width of the image(s) is primarily defined by the detector configuration (the way in which detector elements are combined into data channels along the z -axis). Throughout this report, the total width of the pre-patient collimation will be referred to as the “beam collimation.”

Figure 2 illustrates different detector configurations (along the z -direction) used by MDCT vendors for their 4-channel systems. While the GE detector comprises 16 identically sized detector rows, the Siemens, Philips (previously Marconi), and Toshiba systems utilize variably sized detector-rows along the z -direction. With the variable detector-size design, the beam collimation and detector configuration can be chosen in such a manner as to obtain *veffective* detector widths other than those of the detector rows *per se*. For example, the 4-channel Siemens and Philips systems are capable of acquiring four 1-mm slices (i.e., 4 x 1 mm detector configuration)

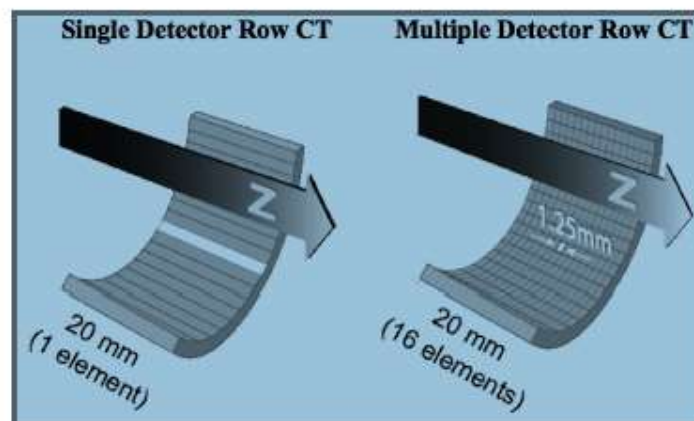


Figure (2-1): The single-detector row CT (SDCT) system on the left has one detector element along the longitudinal axis and many (approx. 900) elements on the arc around the patient.

The width of the detector (relative to the center of the gantry) is 20 mm, although the maximum beam width is only 10 mm. Thus the detector is wider than the x-ray beam. The multiple-detector-row CT (MDCT) system on the right has 16 1.25-mm detector elements along the longitudinal axis for EACH of the approximately 900 positions around the patient. The width of the detector is also 20 mm at its center. The four data

channels allow the acquisition of four simultaneous slices, of either 1.25, 2.5, 3.75, or 5 mm width.

ting the beam collimation to 4 mm. (Throughout this report, the detector configuration will be represented by the product of the number of independent data channels N and the width, along the z -direction at isocenter, imaged by one detector channel T , or $N \times T$ mm). This 4 x 1-mm mode fully irradiates the two central 1-mm detector elements and partially irradiates the two neighboring 1.5-mm rows to effectively give a 4 x 1 mm acquisition. This is accomplished with use of post-patient collimation along the z -axis. Figure 3 details the MDCT detector geometries for 64- channel systems from four major manufacturers

As noted above, MDCT allows information from multiple detector rows to be combined into one data channel. For example, when the 4-channel Toshiba system utilizes its maximum beam collimation (32 mm), four 8-mm virtual detector rows may be formed by combining the signal from eight 1-mm wide detector-rows into a single channel. A significant advantage of MDCT is that signals from multiple data channels may be summed to yield slice widths that are larger than the width corresponding to a given data channel. This may be done retrospectively, allowing, for example, a 4 x 1.25-mm data acquisition to be presented as one 5-mm thick slice, two 2.5-mm slices, four 1.25-mm slices, or all of these options.

When specifying an imaging protocol, it is very important to note the detector configuration used to acquire the desired slice thickness, as this significantly affects the subsequent retrospective reconstruction options (for thinner or thicker images) and the radiation efficiency of the system (i.e., patient dose). For instance, using an MDCT scanner one might acquire 5-mm slices either by using a wide beam collimation (4 x 5 mm) or by utilizing a narrow beam collimation (4 x 1.25 mm). The wide beam collimation allows much faster z -coverage, while the slower narrow beam

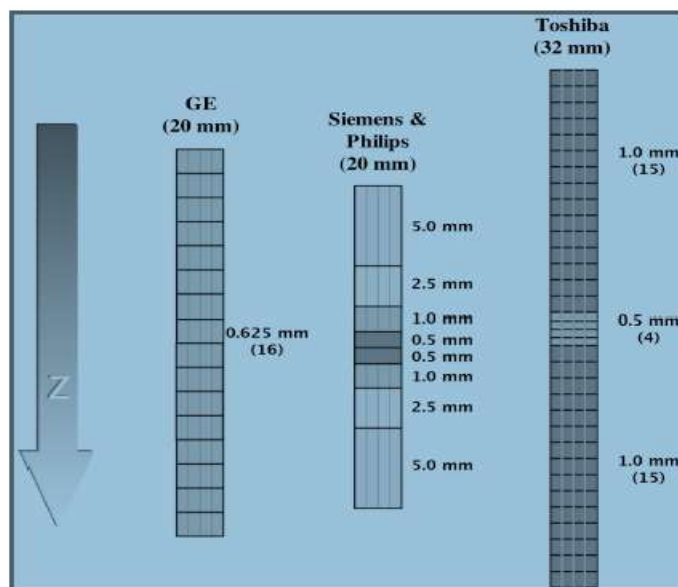


Figure (2-2): Diagram of the detector geometries used in the 4-channel MDCT systems from the four major CT manufacturers.

The detector geometry used on both the Siemens and the Philips (Marconi) 4-channel scanners was co-developed by Siemens and Elscint. In this design, the 20-mm wide detector array uses eight rows of varying widths to allow simultaneous scanning of up to four 5-mm thick slices.

collimation acquisition allows retrospective reconstruction of narrower slice widths. As will be discussed later, this trade-off is complicated by the competing issues of (1) the desire for thin slices, (2) the increase in image noise for thin slices, (3) the relative radiation dose inefficiency of narrow beam collimations, and (4) data management issues (reconstruction and transfer times, archive and filming costs).

The advent of helical CT introduced an additional acquisition parameter into the CT vocabulary, *pitch*. Pitch was defined as the ratio of the table travel per x-ray tube rotation to the slice width (which was typically, but not always, equal to the beam collimation). The advent of MDCT introduced significant confusion regarding the definition of pitch, as some manufacturers used an altered definition of pitch that related the table travel per x-ray tube rotation to the width of an individual data channel. For example, using a 4-channel system ($N_{max} = 4$), a reconstructed slice width of 5 mm, a detector configuration of 4 x 5 mm (nominal beam collimation = 20 mm), and a table travel per rotation of 15 mm, the definition of pitch originally used with helical CT would yield

$15/20 = 0.75$. The manufacturer's altered definition yielded $15/5 = 3$. Hence, the

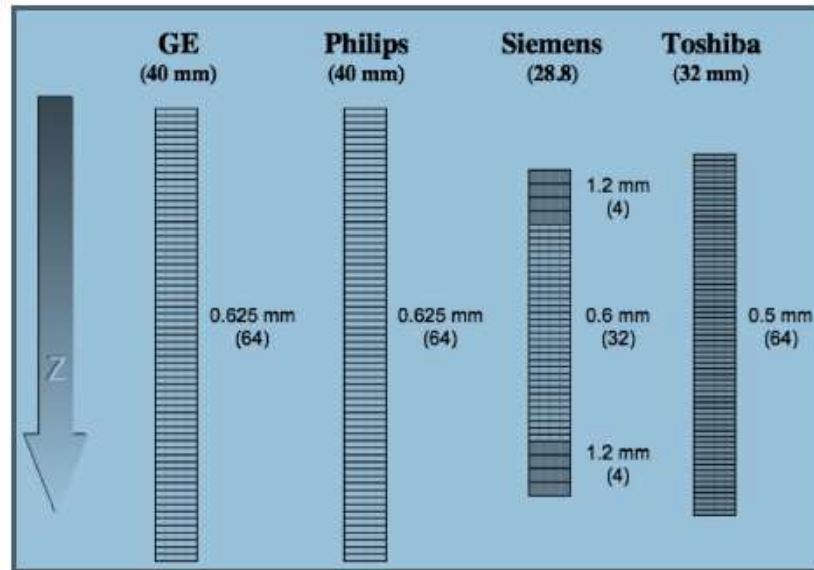


Figure (2-3): Diagram of the detector geometries used in 64-channel MDCT from four major manufacturers. The Siemens 64-MDCT uses 32 submillimeter detectors and a moving focal spot to achieve 64 overlapping slice measurements.

two definitions differed by a factor of 4 (N , the number of data channels used in the acquisition).

As the number of data channels increased, the use of two definitions of pitch caused further confusion, as well as difficulty in comparing scan protocols and radiation dose values. Hence, the IEC reissued their CT safety standard and specifically addressed the definition of pitch, reestablishing the original definition of pitch (table travel normalized to the total beam collimation) as the only acceptable definition of pitch. CT manufacturers altered their user interfaces accordingly for newer software releases, although older scanners with early software versions and the altered definition may still be in use. The IEC definition expresses a concept of pitch that is common

to both SDCT and MDCT. From a radiation dose perspective, it is imperative to use the appropriate pitch definition (table travel per total beam collimation) because it conveys the degree of overlap of the radiation beam: a pitch of 1.0 indicates contiguous radiation beams, a pitch less than 1.0 indicates overlap of the radiation beams, and a pitch greater than 1.0 indicates gaps between the radiation beams. If this

definition of MDCT pitch were not used in a radiation dose calculation, the result would be a factor of N too small

As in SDCT, the tube current and the exposure time (per rotation) govern the number of x-ray photons utilized per rotation, which is given by $\text{mA} \cdot \text{s}$, or simply mAs (milliamperes-second). It is important to note that just as in SDCT, mAs is indicative of *relative* output (radiation exposure) of a CT x-ray tube on a given type of CT scanner, at a given kVp. It does not indicate the absolute output (dose), as the exposure per mAs varies significantly between CT scanner manufacturers,

models, and kVp settings. Thus, 200 mAs/rotation may produce significantly different results (in dose and image quality) on different types of CT scanners and at different kVp settings. For the purpose of comparing radiation dose, mAs should be scaled to a value on each system that gives equivalent image quality (spatial resolution, contrast resolution, and noise). Two manufacturers (Siemens and Philips) report the mAs as the average mAs along the z-axis, called either *effective* mAs or mAs/slice, where effective mAs or mAs/slice is defined as the true mAs/pitch (here they employ the IEC definition of pitch). This distinction between mAs and average mAs along the z -axis is very important, particularly when correcting CT dose metrics for beam overlap or gaps (pitch).

In MDCT, noise is dependent on pitch (this is not true in SDCT). Thus, as pitch is increased, MDCT scanner software may automatically increase the mA such that the image noise (and patient dose) remains relatively constant with changing pitch values. When the *effective* mAs or mAs/slice is used, noise appears to be unaffected by pitch, since noise remains constant as pitch is varied for a constant value of effective mAs or mAs/slice. Thus, the user may be unaware that the actual mA was increased in systems that use the average mAs along the z -axis concept. Another manufacturer (GE) also helps the user to keep image noise constant as pitch is changed. On the GE system, as parameters such as detector configuration, pitch, or image width are changed, the mA value is automatically adjusted to the value that will keep image noise the same. In this scenario, the mA parameter field is flagged (turned orange) to alert the user of the change in the prescribed mA value.

In summary, MDCT technology offers significant improvements in the variety, quality, and speed of CT clinical applications. The technology will continue to change at a rapid pace, and radiologists, technologists, physicists and department administrators will all need to reevaluate existing practice strategies and exam protocols to successfully integrate increasingly complex MDCT scanners into their CT practice. This expected increase in utilization must be accompanied by awareness and understanding of radiation dose issues. In addition, as CT technology develops, the revision or updating of existing definitions, particularly with respect to CT dosimetry, may be required.

The purpose of this report is to provide a reference for the physics community to clarify existing definitions related to CT dosimetry, to describe methods to measure or calculate CT dose descriptors, and to discuss the issues necessary to make clinically relevant decisions regarding CT technique factors and their impact on radiation dose

2.3 Definitions of Quantities For Assessing Dose In CT:

CTDI, CTDIFDA,

CTDI100, CTDIW, CTDIVOL, DLP, E

2.3.1 Computed Tomography Dose Index (CTDI)

The CTDI is the primary dose measurement concept in CT

$$CTDI = \frac{1}{nt} \int_{-\infty}^{\infty} D(z) dz,$$

where

$D(z)$ = the radiation dose profile along the z-axis,

N = the number of tomographic sections imaged in a single axial scan. This is equal to

T = the number of data channels used in a particular scan. The value of N may be less than or equal to the maximum number of data channels available on the system, and the width of the tomographic section along the z -axis imaged by one data channel. In multiple-detector-row

(multislice) CT scanners, several detector elements may be grouped together to form one data channel. In single-detector-row (single-slice)

CT, the z -axis collimation (T) is the nominal scan width.

CTDI represents the average absorbed dose, along the z -axis, from a series of contiguous irradiations. It is measured from one *axial* CT scan (one rotation of the x-ray tube), and is calculated by dividing the integrated absorbed dose by the nominal total beam collimation. The CTDI is always measured in the axial scan mode for a single rotation of the x-ray source, and theoretically estimates the average dose within the central region of a scan volume consisting of multiple, contiguous CT scans [Multiple Scan Average Dose (MSAD)] for the case where the scan length is sufficient for the central dose to approach its asymptotic upper limit. The MSAD represents the average dose over a small interval ($-I/2, I/2$) about the center of the scan length ($z = 0$) for a scan interval I , but requires multiple exposures for its direct measurement. The CTDI offered a more convenient yet nominally equivalent method of estimating this value, and required only a single-scan acquisition, which in the early days of CT, saved a considerable amount of time.

2.3.2 CTDI_{FDA}

Theoretically, the equivalence of the MSAD and the CTDI requires that all contributions from the tails of the radiation dose profile be included in the CTDI dose measurement. The exact integration limits required to meet this criterion depend upon the width of the nominal radiation beam and the scattering medium. To standardize CTDI measurements (infinity is not a likely measurement parameter), the FDA introduced the integration limits of $\pm 7T$, where T represented the nominal slice width²⁶. Interestingly, the original CT scanner, the EMI Mark I, was a dual-detector-row system. Hence, the nominal radiation beam width was equal to twice the nominal slice width (i.e., $N \times T$ mm). To account for this, the CTDI value must be normalized to $1/NT$:

$$CTDI_{FDA} = \frac{1}{NT} \int_{-7T}^{7T} D(z) dz.$$

Unfortunately, the limits of integration were not similarly expressed in terms of NT , allowing for the potential underestimation of the MSAD by the CTDI. For the technology available circa 1984, the use of NT in the integration limits was deemed unnecessary at the time²⁷. The scattering media for CTDI measurements were also standardized by the FDA²⁶. These consist of two polymethylmethacrylate (PMMA, e.g., acrylic or Lucite™) cylinders of 14-cm length. To estimate dose values for head examinations, a diameter of 16 cm is to be used. To estimate dose values for body examination, a diameter of 32 cm is to be used. These are typically referred to, respectively, as the head and body CTDI phantoms.

2.3.3 CTDI₁₀₀

CTDI₁₀₀ represents the accumulated multiple scan dose at the center of a 100-mm scan and underestimates the accumulated dose for longer scan lengths. It is thus smaller than the equilibrium dose or the MSAD. The CTDI₁₀₀, like the CTDI_{FDA}, requires integration of the radiation dose profile from a single axial scan over specific integration limits. In the case of CTDI₁₀₀, the integration limits are ± 50 mm, which corresponds to the 100-mm length of the commercially available “pencil” ionization chamber.

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

The use of a single, consistent integration limit avoided the problem of dose overestimation for narrow slice widths (e.g., <3 mm)²⁴. CTDI₁₀₀ is acquired using a 100-mm long, 3-cc active volume CT “pencil” ionization chamber and the two standard CTDI acrylic phantoms [head (16-cm diameter) and body (32-cm diameter)]. The measurement must be performed with a *stationary* patient table.

The pencil chamber of active length l is not really measuring exposure (X), or air kerma, but rather the integral of the single rotation dose profile $D(z)$. Although the exposure (or air kerma) meter may convert the charge collected into an apparent exposure reading in roentgens (R) (or air kerma reading in milligray [mGy]), the measured value, called the “meter

reading,” actually represents the average exposure (or air kerma) over the chamber length Δz . That is,

$$\text{Meter Reading} = \frac{1}{\Delta z} \int_{-\Delta z/2}^{\Delta z/2} X(z) dz = \frac{1}{f \Delta z} \int_{-\Delta z/2}^{\Delta z/2} D(z) dz$$

where f is the f-factor (exposure-to-dose conversion factor, $D = f \cdot X$). Considering the above definition of CTDI₁₀₀ ($\Delta z = 100$ mm), it is clear that .

$$CTDI = \frac{\int \left(\frac{\text{rad}}{\text{R}}\right) \cdot (\text{mm}) \cdot \text{meter reading (R)}}{N \cdot T(\text{MM})}$$

where:-

C =the unitless chamber calibration factor (typically near 1.0) which is required to

correct the meter reading for temperature and pressure and into true exposure (if the

calibration and measurement beam qualities differ sufficiently to require it).

One must use the f-factor (f) appropriate to the task at hand to convert exposure (R) to

absorbed dose (rad):

- 0.78 rad/R for calculation to dose to acrylic (e.g., CTDIFDA).
- 0.94 rad/R for tissue dose estimates.
- 0.87 rad/R for dose to air and calculation of or comparison to CTDI₁₀₀ or CTDI_w (see section 3.4).
- These values correspond to the typical CT kVp value of 120 kVp, which corresponds to an effective energy of approximately 70 keV.
- For scans at other tube voltage settings, the f-factors must be chosen accordingly.

When an ion chamber measurement is given in air kerma (mGy), care must be taken to indicate which f-factor is used, if any, since the chamber reading and CTDI value are both given in units of mGy:

- 1.06 mGy/mGy for dose to tissue
- 0.90 mGy/mGy for dose to Lucite
- 1.00 mGy/mGy for dose to air.

2.3.4 Weighted CTDI_w

The CTDI varies across the field of view (FOV). For example, for body CT imaging, the CTDI is typically a factor or two higher at the surface than at the center of the FOV. The average CTDI across the FOV is estimated by the Weighted CTDI (CTDI_w), where

The values of 1/3 and 2/3 approximate the relative areas represented by the center and edge values³¹. CTDI_w is a useful indicator of scanner radiation output for a specific kVp and mAs. According to IEC 60601-2-44, CTDI_w must use CTDI₁₀₀ as described above and an f-factor for air (0.87 rad/R or 1.0 mGy/mGy)

2.3.5 Volume CTDI_{vol}

To represent dose for a specific scan protocol, which almost always involves a series of scans, it is essential to take into account any gaps or overlaps between the x-ray beams from consecutive rotations of the x-ray source. This is accomplished with use of a dose descriptor known as the Volume CTDI_w (CTDI_{vol}), where

$$CTDI_{vol} = \frac{N \times T}{I} \times CTDI_w$$

and I = the table increment per axial scan (mm). Since pitch is defined¹⁹ as the ratio of the table travel per rotation (I) to the total nominal beam width ($N \times T$)

$$\text{pitch} = I / (N \times T)$$

Thus, Volume CTDI can be expressed as

$$CTDI_{vol} = 1 / \text{pitch} \times CTDI_w$$

Whereas $CTDI_w$ represents the average absorbed radiation dose over the x and y directions at the center of the scan from a series of axial scans where the scatter tails are negligible beyond the 100-mm integration limit, $CTDI_{vol}$ represents the average absorbed radiation dose over the x , y , and z directions. It is conceptually similar to the MSAD, but is standardized with respect to the integration limits (± 50 mm) and the f -factor used to convert the exposure or air kerma measurement into dose to air.

The $CTDI_{vol}$ provides a single CT dose parameter, based on a directly and easily measured quantity, which represents the average dose within the scan volume for a standardized (CTDI) phantom. The SI units are milligray (mGy). $CTDI_{vol}$ is a useful indicator of the dose to a standardized phantom for a specific exam protocol, because it takes into account protocol-specific information such as pitch. Its value may be displayed prospectively on the console of newer CT scanners, although it may be mislabeled on some systems as $CTDI_w$. The IEC consensus agreement on these definitions is used on most modern scanners.

While $CTDI_{vol}$ *estimates* the average radiation dose within the irradiated volume for an object of similar attenuation to the CTDI phantom, it does not represent the average dose for objects of substantially different size, shape, or attenuation or when the 100-mm integration limits omit a considerable fraction of the scatter tails. Further, it does not indicate the total energy deposited into the scan volume because it is independent of the length of the scan. That is, its value remains unchanged whether the scan coverage is 10 or 100 cm. It estimates the dose for a 100-mm scan length only, even though the actual volume-averaged dose will increase with scan length up to the limiting equilibrium dose value.

2.3.6 Dose-Length Product (DLP)

To better represent the overall energy delivered by a given scan protocol, the absorbed dose can be integrated along the scan length to compute the Dose-Length Product (DLP), where

$$DLP \text{ (mGy-cm)} = CTDI_{vol} \text{ (mGy)} \times \text{scan length (cm)}.$$

The DLP reflects the total energy absorbed (and thus the potential biological effect) attributable to the complete scan acquisition. Thus, an abdomen-only CT exam might have the same $CTDI_{vol}$ as an

abdomen/pelvis CT exam, but the latter exam would have a greater DLP, proportional to the greater z -extent of the scan volume.

In helical CT, data interpolation between two points must be performed for all projection angles. Thus, the images at the very beginning and end of a helical scan require data from z -axis projections beyond the defined “scan” boundaries (i.e., the beginning and end of the anatomic range over which images are desired). This increase in DLP due to the additional rotation(s) required for the helical interpolation algorithm is often referred to as “overranging.” For MDCT scanners, the number of additional rotations is strongly pitch dependent, with a typical increase in irradiation length of 1.5 times the total nominal beam width

The implications of overranging with regard to the DLP depends on the length of the imaged body region. For helical scans that are short relative to the total beam width, the dose efficiency (with regard to overranging) will decrease. For the same anatomic coverage, it is generally more dose efficient to use a single helical scan than multiple helical scans

Table 1 illustrates the differences in CTDI_{vol} and DLP for typical CT exams. The values are meant to be demonstrative only; they can vary by scanner model, vendor, and image quality requirements. Note that a change in technique (mAs/rotation) affects the CTDI_{vol} (and therefore also the DLP), while a change in acquisition length (at the same technique) is only reflected by the DLP.

2.3.7 Limits To CTDI Methods

For body scan lengths of 250 mm or more, the accumulated dose closely approaches the limiting equilibrium dose. However, CTDI₁₀₀ underestimates the equilibrium dose CTDI (or MSAD for pitch of unity) by a factor of approximately 0.6 on the central axis and by about 0.8 on the periph-

Table (2-1) : Illustrative values for CTDI_{vol} and DLP for common CT exams for (a) 4-channel MDCT and (B) 16-channel MDCT

Table 1a:4- channel MDCT(120KVp)

<i>Exam</i>	<i>Beam Collimation</i>	<i>Pitch</i>	<i>mAs per Rotation</i>	<i>Scan Length (cm)</i>	<i>CTDI_{vol} (mGy)</i>	<i>DLP (mGy-cm)</i>
Head	4 x 2.5	Axial	250	15	55.0	825
Chest	4 x 5	0.75	100	40	12.0	480
Abdomen	4 x 5	0.75	150	20	19.1	382
Abdomen & Pelvis ≈	4 x 5	0.75	150	40	19.1	764

Table 1b: 16-channel MDCT (120 kVp)

<i>Exam</i>	<i>Beam Collimation</i>	<i>Pitch</i>	<i>mAs per Rotation</i>	<i>Scan Length (cm)</i>	<i>CTDI_{vol} (mGy)</i>	<i>DLP (mGy-cm)</i>
Chest	16 x 1.25	0.983	150	35	13.3	466
Abdomen	16 x 1.25	0.983	212	28	18.8	526
Pelvis	16 x 1.25	0.983	212	25	18.8	470

The total energy imparted is underestimated by the DLP by a factor of about 0.7 for all scan lengths. In order to measure the equilibrium dose, a body phantom length of almost 400 mm is required. Since a pencil chamber of this length is not practical, direct measurement of the MSAD using a conventional ion chamber can be utilized. Such a method can be utilized to emulate a “virtual” pencil chamber of arbitrary length up to the available phantom length.

2.3.8 Effective Dose (E)

It is important to recognize that the potential biological effects from radiation depend not only on the radiation dose to a tissue or organ, but also on the biological sensitivity of the tissue or organ irradiated. A 100-mGy dose to an extremity would not have the same potential biological effect (detriment) as a 100-mGy dose to the pelvis³⁵. Effective dose, *E*,

is a dose descriptor that reflects this difference in biologic sensitivity. It is a single dose parameter that reflects the risk of a nonuniform exposure in terms of an equivalent whole-body exposure. The units of effective dose are sieverts (usually millisieverts (mSv) are used in diagnostic radiology).

The concept of effective dose was designed for radiation protection of occupationally exposed personnel. It reflects radiation detriment averaged over gender and age, and its application has limitations when applied to medical populations. However, it does facilitate the comparison of biologic effect between diagnostic exams of different types. The use of effective dose facilitates communication with patients regarding the potential harm of a medical exam that uses ionizing radiation. For example, when a patient inquires, “What dose will I receive from this exam?” an answer in the units of mGy or mGy-cm will not likely answer the more fundamental, but perhaps unspoken, question, “What is the likelihood that I will be harmed from this exam?” Characterizing the radiation dose in terms of effective dose and comparing that value to other radiation risks, for instance one year’s effective dose from naturally occurring background radiation, better conveys to the patient the relative potential for harm from the medical exam. Table 2 provides typical values of effective dose for several common imaging exams (CT and non-CT), as well as the annual level of naturally occurring background radiation in the United States (3.0 mSv).

Table (2-2) Type effective dose values for several common imaging exam (CT and non-CT)

<i>Non-CT Typical Effective Dose Values (mSv)</i>		<i>CT Typical Effective Dose Values (mSv)</i>	
Hand radiograph	<0.1	Head CT	1-2
Dental bitewing	<0.1	Chest CT	5-7
Chest radiograph	0.1-0.2	Abdomen CT	5-7
Mammogram	0.3–0.6	Pelvis CT	3-4
Lumbar spine radiograph	0.5–1.5	Abdomen & pelvis CT	8-14
Barium enema exam	3-6	Coronary artery calcium CT	1-3
Coronary angiogram (diagnostic)	5-10	Coronary CT angiography	5-15
Sestamibi myocardial perfusion	13-16		
Thallium myocardial perfusion	35-40		

Note: Average U.S. background radiation from naturally occurring sources ≈ 3.0 mSv (range 1–10 mSv)⁶⁷

It is important to remember, however, that the effective dose describes the relative “wholebody” dose for a particular exam and scanner, but is *not* the dose for any one individual. Effective dose calculations use many assumptions, including a mathematical model of a “standard” human body that does not accurately reflect any one individual (it is androgynous and of an age representative of a radiation worker). Effective dose is best used to optimize exams and to compare risks between proposed exams. It is a broad measure of risk, and as such, should not be quoted with more than one or two significant digits.

The most direct way of estimating doses to patients undergoing CT examinations is to measure organ doses in patient-like phantoms. Another way of obtaining the pattern of energy deposition in patients undergoing CT examinations is by calculation. Computations that use Monte Carlo methods follow the paths of a large number of x-rays as they interact with a virtual phantom and estimate the probability of the dominant interaction processes (i.e., Compton scatter and photoelectric absorption). This type of calculation assumes that the patient resembles the phantom used for measurements or Monte Carlo simulation. When patients differ in size and composition, appropriate corrections might need to be used. The resultant information is the absorbed dose to a specified tissue, which may be used to predict the biological consequences to that (single) tissue. CT examinations, however, irradiate multiple tissues having different radiation sensitivities. The effective dose takes into account how much radiation is received by an individual tissue, as well as the tissue’s relative radiation sensitivity.

Specific values of effective dose can be calculated using several different software packages, which are based on the use of data from one of two sources, the National Radiological Protection Board (NRPB) in the United Kingdom or the Institute of Radiation Protection (GSF) in Germany. A free Excel spreadsheet can be downloaded from www.impactscan.org to perform organ dose and effective dose estimates using the NRPB organ dose coefficients. Other packages are available for purchase.

To minimize controversy over differences in effective dose values that are purely the result of calculation methodology and data sources, a generic estimation method was proposed by the European Working Group for Guidelines on Quality Criteria in Computed Tomography²¹. Effective

dose values calculated from the NRPB Monte Carlo organ coefficients were compared to DLP values for the corresponding clinical exams to determine a set of coefficients k , where the values of k are dependent only on the region of the body being scanned (head, neck, thorax, abdomen, or pelvis) (Table 3). Using this methodology, E can be estimated from the DLP, which is reported on most CT systems:

Table (2-3): Normalized effective dose per dose-length product (DLP) for adults (standard physique) and pediatric patients of various ages over various body regions. Conversion factor for adult head and neck and pediatric patients assume use of the head CT dose phantom (16 cm). All other conversion factors assume use of the 32-cm diameter CT body phantom

Region o Body	K(mSv mGy⁻¹ cm⁻¹)				
	<i>0 year old</i>	<i>1 year old</i>	<i>5 year old</i>	<i>10 year old</i>	<i>Adult</i>
<i>Exam</i>					
Head and neck	0.013	0.0085	0.0057	0.0042	0.0031
Head	0.011	0.0067	0.0040	0.0032	0.0021
Neck	0.017	0.012	0.011	0.0079	0.0059
Chest	0.039	0.026	0.018	0.013	0.014
Abdomen ~ & Pelvis	0.049	0.030	0.020	0.015	0.015
Trunk	0.0440.039	0.028	0.019	0.014	0.015

$$E \text{ (mSv)} = k \times DLP.$$

The values of E predicted by DLP and the values of E estimated using more rigorous calculations methods are remarkably consistent, with a maximum deviation from the mean of approximately 10% to 15%⁴¹. Hence, the use of DLP to estimate E appears to be a reasonably robust method for estimating effective dose. Similarly, Huda has compared effective dose, as calculated from the NRPB data, to estimates of energy imparted in order to develop conversion coefficients by which to later estimate effective dose from energy imparted.

2.4 Overview of Methods For Dose Reduction In Ct

Recently several new approaches have focused on reducing the radiation dose required to create a CT image of appropriate diagnostic quality. Current dose reduction technical initiatives by researchers and manufacturers can be placed into one of the following general categories.

2.4.1 X-ray Beam Filtration

The use of an absorbent material between the x-ray tube and the patient can be used to “harden” the beam such that low-energy x-rays (which contribute disproportionately to absorbed dose) are reduced, or to “shape” the x-ray beam to deliver dose in the most appropriate spatial distribution.

Previously, only head and body beam shaping (e.g., “bowtie”) filters were available. Recently, manufacturers have added filters more specific to cardiac imaging or different-sized patients.

2.4.2 X-ray Beam Collimation

The use of a very attenuating material between the x-ray tube and the patient should be used to limit the x-ray beam to the minimal dimensions required. Such collimation occurs along the z -axis to define the radiation beam width. Additional collimation after the patient to further define the image width, whether an absorbent material or electronic, causes radiation dose to the patient to be wasted. Finally, the fan angle of the beam should be collimated to the diameter of the patient to reduce the amount of bypass that can then be scattered back towards the patient or towards personnel. Such inplane beam collimation is typically implemented by use of an appropriate scan FOV (shaping filter).

MDCT Dose Inefficiency At Narrow Beam Collimations

MDCT systems have been observed to have a radiation dose inefficiency at narrow beam collimations, resulting in a higher CTDI for the narrow beam collimations required for narrow slice widths. In SDCT, CTDI is generally independent of slice width (although for some SDCT systems, the CTDI can increase by as much as a factor of 2 for scan widths less than 2 mm).

The dose inefficiency in current MDCT designs is due to unused x-ray beam that strikes outside of the active area of the detector (along the z -direction). The z extent of this unused portion of the x-ray beam is

approximately constant in size for the various detector configurations; thus the inefficiency caused by the unused radiation is relatively greater at narrow beam collimations. In 4-channel MDCT systems, the narrow beam dose inefficiency can be substantial, resulting in

as much as a 40% to 50% dose increase for the narrow beam collimations (4x1 mm or 4x1.25 mm) relative to the widest beam collimations (4x5 mm or 4x8 mm). For submillimeter beam collimations on 4-channel MDCT systems, this dose increase can be over 100% relative to the widest beam collimations. The use of a greater number of data channels (16 or more) covering larger z -axis extents of the detector increases the dose efficiency of MDCT to nearly that of SDCT.

2.4.3 X-ray Tube Current (mAs) Modulation and Automatic Exposure Control (AEC):

It is technologically feasible for CT systems to adjust the x-ray tube current (mA) in real-time during gantry rotation in response to variations in x-ray intensity at the detector, much as fluoroscopic x-ray systems adjust exposure automatically. This capability, in various implementations, is available commercially on MDCT systems in response to wide interest from the radiology community. Some systems adapt the tube current based on changes in attenuation along the z -axis, others adapt to changes in attenuation as the x-ray tube travels around the patient. The ideal is to combine both approaches with an algorithm that “chooses” the correct tube current to achieve a predetermined level of image noise.

By decreasing or increasing the x-ray tube current, the radiation output of the tube is proportionately changed. Image noise is dominated by the noisiest projection (which corresponds to the most attenuating paths through the patient). Hence data acquired through body parts having less attenuation can be acquired with substantially less radiation without negatively affecting the final image noise. This principle can be applied to modulate the mA angularly about the patient (anteriorposterior [AP] vs. lateral) as well as along the z -axis (neck vs. shoulders); the tube current can also be modulated within the cardiac cycle (systole vs. diastole), or with respect to sensitive organs (PA vs. AP).

With regard to cardiac CT, the radiation dose for a retrospectively gated exam, where the x-ray tube is kept continuously on throughout the

acquisition, can be dramatically decreased if the tube current is reduced during portions of the cardiac cycle that are not likely to be of interest for the reconstructed images. Thus, in addition to modulation of the tube current based on patient attenuation, the tube current can be modulated by the ECG signal. Since cardiac motion is least during diastole and greatest during systole, the projection data are least likely to be corrupted by motion artifact for diastolic-phase reconstructions. Accordingly, the tube current is reduced during systole. Dose reductions of approximately 50% have been reported using such a strategy. The implementation of these and other dose reduction strategies is expected industrywide over the next several years, in response to the strong concern about the radiation dose from CT from both the public at large and the medical community

In addition to technical methods of dose reduction, investigators are working to determine clinically acceptable levels of image noise for a variety of diagnostic tasks. That is, high-contrast exams

(e.g., lung, skeletal, colon, sinus) require much less dose (can tolerate higher noise levels) compared to low-contrast exams (e.g., brain, liver, and other abdominal organs). If the required noise level can be predefined, CT systems can use technical approaches to deliver the minimum dose required to achieve the specified noise level. The definition of a robust and standardized noise metric is required, however, to allow a manufacturer-independent method of prescribing the desired image quality.

2.4.4 Size-or Weight-based Technique Charts

Unlike traditional radiographic imaging, a CT image never looks “overexposed” in the sense of being too dark or too light; the normalized nature of CT data (i.e., CT numbers represent a fixed amount of attenuation relative to water) ensures that the image always appears properly exposed. As a consequence, CT users are not technically compelled to decrease the tube-current-time product (mAs) for small patients, which may result in excess radiation dose for these patients. It is, however,

a fundamental responsibility of the CT operator to take patient size into account when selecting the parameters that affect radiation dose, the most basic of which is the mAs.

As with radiographic and fluoroscopic imaging, the operator should be provided with appropriate guidelines for mAs selection as a function of patient size. These are often referred to as technique charts. While the tube current, exposure time, and tube potential can all be altered to give the appropriate exposure to the patient, in CT users most commonly standardize the tube potential (kVp) and gantry rotation time (s) for a given clinical application. The fastest rotation time should typically be

used to minimize motion blurring and artifact, and the lowest kVp consistent with the patient size should be selected to maximize image contrast.

Although scan parameters can be adapted to patient size to reduce radiation dose, it is important to remember certain caveats when contemplating such adjustments. First, body regions such as the head do not vary much in size in the normal population, so modification of scan parameters may not be applicable here based on head size.

Numerous investigators have shown that the manner in which mA should be adjusted as a function of patient size should be related to the overall attenuation, or thickness, of the anatomy of interest as opposed to patient weight, which is correlated to patient girth, but not a perfect surrogate as a function of anatomic region. The exception is for imaging of the head, where attenuation is relatively well defined by age, since the primary attenuation comes from the skull and the process of bone formation in the skull is age dependent.

Clinical evaluations of mA-adjusted images have demonstrated that radiologists do not find the same noise level acceptable in small patients as in larger patients⁵⁹. Because of the absence of adipose tissue between organs and tissue planes, and the smaller anatomic dimensions, radiologists

tend to demand lower noise images in children and small adults relative to larger patients. For body CT imaging, typically a reduction in mA (or mAs) of a factor of 4 to 5 from adult techniques is acceptable in infants. For obese patients, an increase of a factor of 2 is appropriate. For head CT imaging, the mAs reduction from an adult to a newborn of approximately a factor of 2 to 2.5 is appropriate. Sample technique charts are provided in

appendix A. To achieve increased exposure for obese patients, either the rotation time, or the tube potential, may also need to be increased

2.4.5 Detector Geometric Efficiency

Ideally, all of the photons that pass through the patient should be used in the image formation process. However, the conversion of photon energy to electrical signal is not a 100% efficient process (although it exceeds 90% for modern scintillating detectors). New detector materials having even

higher absorption and conversion efficiencies are of course desirable, as are detector and signal processing electronics with very low inherent noise levels. Additionally, the small detector elements are divided along the detector arc and along the z axis with radiation-absorbing septa (walls). These septa also provide essential optical isolation between detector elements, but they, along with the very fine signal transmission wires, create “dead spaces” in the detector and hence waste radiation dose. As detectors continue to be divided into smaller and smaller discrete elements, the geometric efficiency of the detector systems must be maintained. One important step in reducing the dead space has been to attach and route the signal transmission wires underneath each detector element, instead of between detector elements. However, ongoing reductions in voxel size will likely be limited by the exponential increase in image noise that would accompany such changes

2.4.6 Noise Reduction Algorithms

Data processing can be performed on the raw data (in sinogram space) or on already reconstructed images to reduce image noise. A variety of approaches are possible, all of which seek to smooth out random pixel variations (noise) while preserving fine detail and structure (signal). With a successful noise reduction scheme, an image of adequate quality can be acquired with a reduced patient dose

2.5 .Clinical Utility of Ctdivol

The use of routinely displayed scan parameters such as mAs and kVp is minimally successful in predicting dose⁶³. Rather than relying on parameters such as mAs, kVp, and pitch, the use of CTDIvol provides a single “dose metric” by which users can benchmark the prescribed output for a given exam against national averages, already having the effects of

pitch, detector collimation, xray tube to isocenter distance, and other technical parameters all taken into account⁶⁴. The values of CTDI_{vol} displayed on the user console prior to scan initiation can be compared to published values, such as reference values provided by the American College of Radiology (ACR) and

AAPM, and results of national surveys, such as the Nationwide Evaluation of X-ray Trends (NEXT) study conducted by the Center for Devices and Radiological Health (CDRH). Users prescribing doses above references values should have an appropriate justification .

2.6 Appropriate Use of Ct Dose Values And Risk Parameters:

Effective dose estimates will vary somewhat according to the model of equipment and imaging parameters used. Typical values are given in Table 2. Effective dose estimates are only valid for prospective radiological protection purposes and should not be used for retrospective dose assessments or the detailed estimation of a specific individual's risk. Effective dose can be of some value for comparing doses from different diagnostic and therapeutic procedures and for comparing the radiation risks for different technologies, hospitals, or countries. For risk-benefit assessments for any individual, however, the absorbed dose to irradiated tissues is the more appropriate quantity. Patient effective doses may also be compared with background radiation exposures from natural sources, which in the United States averages 3 mSv per year. This allows patients and their families to put medical doses into context and better understand that radiation exposure is an everyday occurrence, not something out of the ordinary. It also obviates the need to convert doses into unfamiliar (and uncertain) radiation risk values.

The most definitive data on the ability of ionizing radiation to induce cancer is obtained from the atomic-bomb survivors cohort, albeit our knowledge of radiation risks at the relatively low radiation dose levels associated with CT scanning is subject to large uncertainties. Current radiation risk estimates are based on a linear no threshold model, which is a topic of ongoing scientific debates.

Radiation risks may also be compared with those encountered in everyday life, such as the risks of dying when smoking cigarettes or the risk of dying in an automobile accident. For patients over 60 years of age, an effective dose of 17 mSv, typical of a cardiac CT angiogram, may be

estimated to have a risk that is comparable to the risk of dying from lung cancer after smoking ~300 packs of cigarettes or the risk of dying in an automobile accident when driving a distance of ~12,000 miles. Although all of these risk estimates are very crude, they do help put radiation risks into a context. More precise risk estimates require taking into account specific organ doses, age, and genders. The assumption that a CT examination has a (small) radiation risk requires that all such exposures need justification and that patient doses need to be kept as low as reasonably achievable (ALARA). Examinations should not be performed when the anticipated patient benefit would be lower than the corresponding patient risk. Further, patients should not be exposed to radiation levels above those required for producing an image of diagnostic quality. A good example of minimizing patient doses is to ensure that the radiographic technique (i.e., mAs setting) is no higher than required to keep the radiographic mottle to an acceptable level. Good practice also requires the use of patient-size-specific protocols and techniques that minimize dose without adversely affecting diagnostic performance. In the United States there are no dose limits for patients undergoing CT examinations. What is deemed to be an acceptable patient dose relies on the professional judgment of the physician in charge of the diagnostic procedure. To this end, it is important to ensure that imaging protocols are continually reviewed such that the choice of radiographic techniques is consistent with ALARA principles. This is best accomplished with the assistance of a diagnostic medical physicist.

2.7 Summary

Modern CT scanners provide two dose parameters that both became available by the scanner manufacturers around 2001: the Volume CTDI (CTDI_{vol}) measured in mGy, and the dose-length product (DLP) measured in mGy-cm. CTDI_{vol} is a measure of the average dose within the scan volume to a standardized phantom. The total amount of radiation delivered to a standardized phantom is represented by the DLP, which is the product of CTDI_{vol} and the scan length. Organ doses in CT are well below the threshold for the induction of deterministic effects (e.g., erythema, epilation). Patient radiation risks in CT are therefore those related to carcinogenesis. An estimate of effective dose (E), which is related to the carcinogenic risk, may be obtained by use of E/DLP conversion factors (Table 3). Effective doses from CT are much higher than effective doses in conventional radiography, but

comparable to those associated with interventional fluoroscopic, diagnostic coronary catheterization, or nuclear medicine examinations. Although this risk from a CT examination is small, it is not zero. Hence, CT examinations should be performed only when a net patient benefit is anticipated. Further, the amount of radiation used should always be kept as low as reasonably achievable (ALARA).

CHAPTER THREE
MATERIAL AND METHODS

CHAPTER THREE

MATERIAL AND METHODS

3.1 Material

Study sample :-

The data used in this study was collected from central medical in Khartoum state .

The data collected from January 2019 was conducted on 50 adult patient during ct examination

There were one machine used in this study G E optima 60 slices

3.2 Methods

The data were collected for patient during the routine CT imaging in this department .

In genral the imaging protocols were used the following setup :

The technologist asked the patient before the exam to conform that symptom match the indication for the CT exam

Shoes exam Protocols which a dresses the Clinic question while maiming does setup patient carefully . Asymmetric positioning can result in decreased image quality an increase in patient dose

3.2 .1 Data collected

The data collected was designed to Assessment the adult patient the collected data in accorded patient sex ,age , tubevoltge and tube current , sclice type . in addition we also record all the Parameters , as will as the C T dose description CT does index volume (milligray) and does length product in (milligray- sent meter) . all the factors that have direct influence on radiation does

3.2.2 Data analysis :

We use Excel Microsoft office to analysis the data

One C T machine were used to collect data during this study

In this study was obtained a measurement of does . We used of the calibration electrometer machine .

In this study were used normalized CTDL values Published by the impact calculate $CTDL_{vol}$ and DLP.

Found the effective does by equation :

$$ED = DLP \times \text{coefficient } K$$

K : coefficient of adult chest CT

CHAPTER FOUR

Results

CHAPTER FOUR

Results

4.1 Results :

The following tables and figures represent the data obtained from 50 patient under were CT examinations for chest with digital imaging in modern medical center in Khartoum state

Table (4-1): sex					
		Frequency	Percent	Valid Percent	Cumulative Percent
Valid	male	22	44.0	44.0	44.0
	female	28	56.0	56.0	100.0
	Total	50	100.0	100.0	

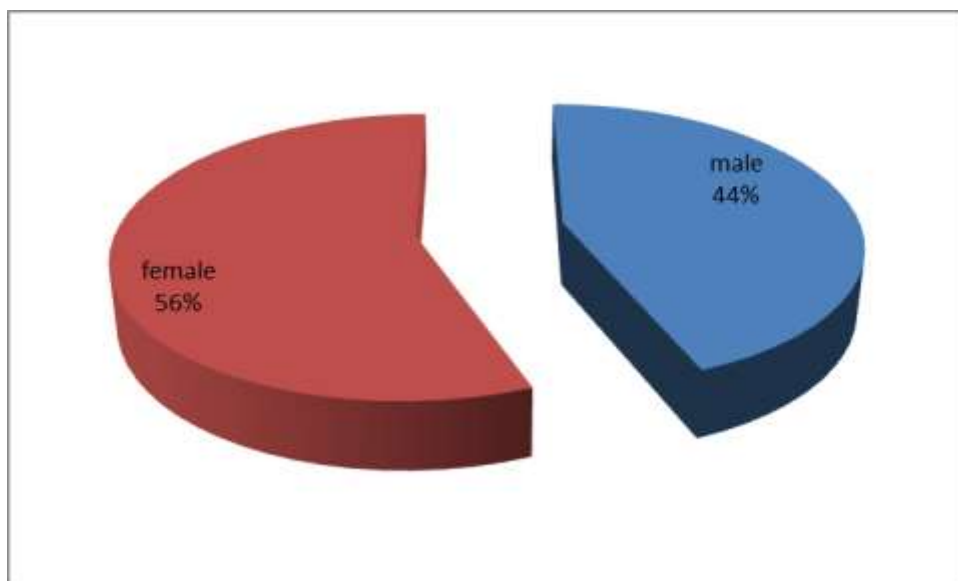


Figure (4-1): show that the rate of female 56% either male 44%

Table (4-2) : age					
		Frequency	Percent	Valid Percent	Cumulative Percent
Valid	less 20 year	5	10.0	10.0	10.0
	20-30 year	4	8.0	8.0	18.0
	30-50 year	7	14.0	14.0	32.0
	50-60 year	6	12.0	12.0	44.0
	more than 60 year	28	56.0	56.0	100.0
	Total	50	100.0	100.0	

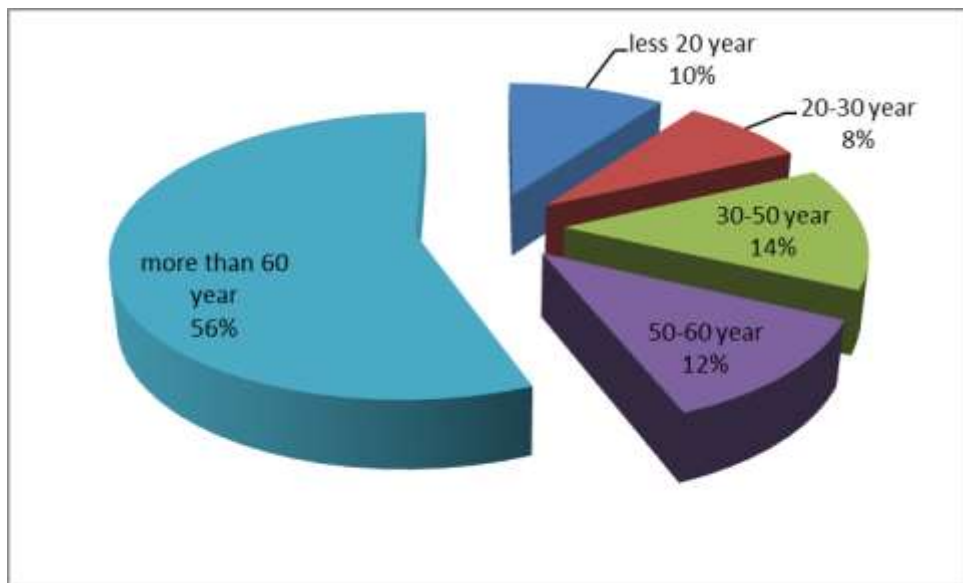


Figure (4-2): show thate rate of ages in this research it is reach more than 60 year 56% and less20 year 10% either(30- 50)reach 14% and (50 - 60) 12%

Table (4-3): Descriptive Statistics summaries the characterizes performance for the CT system and console displaed form the CTDLvol-mGY/cm scanner DLPmGY/cm model acquilion

	N	Minimum	Maximum
sex	50	1	2
age	50	1	5
kv	50	120	120
current Mm	50	10	10
CTDLvol-mGY/cm	50	4	27
DLPmGY/cm	50	52	2066
EMS.V	50	2	29

Table (4-4):Descriptive Statistics show the estimation meam CTDLvol-mGY/cm and DLPmGY/cm and effective dose calculated by CTDLvol-mGY/cm version 4.620 software were used data collection CT scanner DLPmGY/cm

	N	Minimum	Maximum	Mean	Std. Deviation
sex	50	1	2	1.56	.501
age	50	1	5	3.96	1.399
kv	50	120	120	120.00	.000
current Mm	50	10	10	10.00	.000
CTDLvol-mGY/cm	50	4	27	9.13	4.620
DLPmGY/cm	50	52	2066	774.76	498.688
EMS.V	50	2	29	11.29	6.778

Correlations

sex	50	1	2	1.56	.501
age	50	1	5	3.96	1.399
Valid N (listwise)	50				

Case Processing Summary

	Cases					
	Valid		Missing		Total	
	N	Percent	N	Percent	N	Percent
sex * age	50	100.0 %	0	0.0%	50	100.0 %

sex * age Cross tabulation

Count

	age					Total
	less 20 year	20-30 year	30-50 year	50-60 year	more than 60 year	
sex male	3	3	4	1	11	22
sex female	2	1	3	5	17	28
Total	5	4	7	6	28	50

Case Processing Summary

	Cases					
	Included		Excluded		Total	
	N	Percent	N	Percent	N	Percent
sex * age	50	100.0 %	0	0.0%	50	100.0%

ANOVA Table							
			Sum of Squares	df	Mean Square	F	Sig.
sex * age	Between Groups	(Combined)	1.144	4	.286	1.151	.345
	Within Groups		11.176	45	.248		
	Total		12.320	49			

Through the previous tables, it is clear that there is a statistically significant relationship between the exposure of patients or workers in the radiation field in terms of gender, which means both parties, male or female, have the same vulnerability or influence.

Correlations			
		sex	EMS.V
sex	Pearson Correlation	1	.133
	Sig. (2-tailed)		.357
	N	50	50
EMS.V	Pearson Correlation	.133	1
	Sig. (2-tailed)	.357	
	N	50	50

Through the previous tables, it is clear that there is a statistically significant relationship between the exposure of patients or workers in the radiation field in terms of gender, which means both parties, male or female, have the same vulnerability or influence. That is, there are no differences between sex and EMS.V

Correlations			
		EMS.V	age
EMS.V	Pearson Correlation	1	-.171-
	Sig. (2-tailed)		.236
	N	50	50
age	Pearson Correlation	-.171-	1
	Sig. (2-tailed)	.236	
	N	50	50

Correlations					
			EMS.V	sex	
EMS.V	Pearson Correlation		1	.133	
	Sig. (2-tailed)			.357	
	Sum of Squares and Cross-products		2251.098	22.170	
	Covariance		45.941	.452	
	N		50	50	
	Bootstra p ^c	Bias		0	-.002-
		Std. Error		0	.143
		95% Confidence Interval	Lower	1	-.150-
Upper			1	.403	
sex	Pearson Correlation		.133	1	
	Sig. (2-tailed)		.357		
	Sum of Squares and Cross-products		22.170	12.320	
	Covariance		.452	.251	
	N		50	50	
	Bootstra p ^c	Bias		-.002-	0
		Std. Error		.143	0
		95% Confidence Interval	Lower	-.150-	1
Upper			.403	1	
*. Correlation is significant at the 0.05 level (2-tailed).					
**. Correlation is significant at the 0.01 level (2-tailed).					
c. Unless otherwise noted, bootstrap results are based on 1000 bootstrap samples					

CHAPTER FIVE
DISCUSSION ,CONCLUSION
AND RECOMMENDATION

CHAPTER FIVE

DISCUSSION ,CONCLUSION AND RECOMMENDATION

5.1 Discussion :-

This study is intended to assess radiation dose and estimate the radiation entrance skin doses examination of radiation by computerized tomography the data were from a modern medical center .

In table (4) estimation mean effect dose is 11.29 msv and standard deviation 6.778 msv , the maximum dose 29 msv while the minimum 2msv and . It is the estimated entrance surface dose examination for all the examination MGY/MAS to the effective dose MGY/MAS when compared with previous study.

Previous study it is the CT scan can give radiation dose equivalent to 50-450 pairs (posterior anterior and lateral views) of chest radiograph , depending on the CT scan protocol being utilized (10) effective radiation dose equivalent for chest radiograph in two views ranges from 0.06 to 0.25 milli-sieverts (mSv) corresponding dose with CT using conventional examination parameters are 3-27(mSv) and 0.3-0.55 mSv using low radiation CT setting.

In this study the value radiation dose little increase compared with previous study because I do not but may be change size between male and female the patient or CT machine not need to calibration .

The international commission of radiological protection (ICRP) in a publication from 1990 suggested that low level of radiation exposure. The risk of radiation- induced cancer is estimated to be higher in infants and children and lower in the elderly . The scientific basis for many of these projections is weak and has been extrapolated from studies of the effects of higher radiation exposure (gamma rays from atomic explosion), which are greater than dose received in diagnostic radiography .The estimation of risk associated with radiation dose assumes a linear relationship exist between radiation and subsequent risk of development of cancer.

5.2 conclusion :-

The Radiation dose was measured modern medical center using CT GE optima 60 slices .

The higher dose is 29 mSv while the lower dose is 2mSv.

Radiation dose procedures varies from patient to patient.

The main dose variations in the some clinic could be attributed to the techniques , which justify . the important of use radiation dose optimization technologists training does to reducing must be well understood and used .

5.3 Recommendation :-

Using the best strategies available for reducing radiation does to allow radiation in relation to the patients size and weight ,adopted tube current . the radiology community that does must be a slow reasonable achievable (ALARA) While be maintaining diagnostic quality .

The use of CT must also be justified for the diagnostic task.

- i. Must be used tools properly to achieve safe and effective result . Numerous CT protocol optimization strategies have been outlined .
- ii. Implementation high quality CT protocol with radiation exposure appropriate for the clinical setting and size of patient.
- iii. Implementation primary engagement from radiologist ideally in collaboration with CT manufacturers CT technologist and medical physics.

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APPENDICES

Appendix

No	sex	AGE	KV	Current Mm	CTDL _{vol} mGY cph	DLP mGY cm	E MS.V
1.	male	more than 60 year	120	10	9	636	9
2.	female	more than 60 year	120	10	11	1046	15
3.	female	30-50 year	120	10	11	355	5
4.	male	30-50 year	120	10	9	303	4
5.	female	50-60 year	120	10	4	332	4
6.	male	30-50 year	120	10	4	270	4
7.	female	more than 60 year	120	10	9	1076	15
8.	female	50-60 year	120	10	6	345	20
9.	male	more than 60 year	120	10	5	559	8
10.	male	less 20 year	120	10	5	642	9
11.	female	50-60 year	120	10	4	209	3
12.	male	20-30 year	120	10	10	550	8
13.	male	more than 60 year	120	10	6	460	6
14.	female	20-30 year	120	10	4	525	7
15.	female	more than 60 year	120	10	7	1327	19
16.	female	50-60 year	120	10	12	1473	21
17.	female	more than 60 year	120	10	13	722	10
18.	female	more than 60 year	120	10	13	539	8
19.	female	more than 60 year	120	10	6	431	6
20.	female	more than 60 year	120	10	15	956	13
21.	male	20-30 year	120	10	6	2066	29
22.	male	more than 60 year	120	10	9	214	3
23.	female	less 20 year	120	10	7	1057	15
24.	male	20-30 year	120	10	8	1080	15
25.	female	more than 60 year	120	10	9	1819	25
26.	female	more than 60 year	120	10	14	163	2
27.	female	more than 60 year	120	10	5	494	7
28.	female	less 20 year	120	10	6	1959	27
29.	female	more than 60 year	120	10	10	575	8
30.	male	less 20 year	120	10	27	678	9
31.	female	30-50 year	120	10	19	1805	25
32.	male	30-50 year	120	10	12	426	6
33.	male	more than 60 year	120	10	15	778	11
34.	female	more than 60 year	120	10	11	241	3
35.	female	30-50 year	120	10	5	713	10

36.	female	more than 60 year	120	10	11	485	7
37.	male	more than 60 year	120	10	15	437	6
38.	male	less 20 year	120	10	7	599	8
39.	male	30-50 year	120	10	8	965	14
40.	male	more than 60 year	120	10	5	672	10
41.	male	more than 60 year	120	10	15	1079	15
42.	male	more than 60 year	120	10	4	159	13
43.	female	50-60 year	120	10	13	912	10
44.	female	more than 60 year	120	10	11	727	16
45.	female	more than 60 year	120	10	6	1157	7
46.	male	50-60 year	120	10	4	52	20
47.	male	more than 60 year	120	10	5	486	2
48.	female	more than 60 year	120	10	15	1416	17
49.	female	more than 60 year	120	10	5	1568	13
50.	male	more than 60 year	120	10	7	1206	17