

INTEGRATION OF AN INFORMATION SYSTEM TO SUGGEST A DRL MODEL IN RADIOLOGY DEPARTMENT AT AZADI HOSPITAL IN DUHOK-KURDISTAN REGION, IRAQ

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ABSTRACT

Computer Tomography is a useful technique in diagnostic radiology. However, it emits high radiation doses which may cause harm to patients. For this, a quality management system, Diagnostic Reference Levels (DRL) is used in European countries. The aim of this study was to suggest a DRL system in Duhok Hospitals. Patient's data (70 – 80) were collected from Azadi Hospital for three main protocols head, chest and abdomen. Only 15 patients had complete data for calculations for the three protocols. The DRL results were then compared with the ones reported in several European countries. For example, in the head protocol for the a CTDI value of 81, the results of the calculated DLP and ED values were 1504 and 3.16 respectively. These values were found to be much higher than the corresponding values reported in Italy 13.12 and 2.76 respectively. As a result, the DRL using 75th percentiles corresponding to these three dosimetric values were also high. Similar high results for the abdomen protocol were recorded, however, the results of the chest protocol showed to be within the average range. Part two of this work involved designing a Flow Chart based on an Information System to help running the CT scan procedure more effectively. Additionally, an extra repository system was added to this model to solve the short capacity issue in Duhok Hospitals.

KEYWORDS: Data Management Systems, Data Repository, Diagnostic Reference Level, Dose Management, Information Technology, Radiology Information System

1. INTRODUCTION

One of the great challenges that may face today's Hospitals in many countries is the issue of gathering, storing, tracing and maintaining data in regard to patient information (Kruse et. al., 2016). Due to the advancements in Information Technology (IT) several techniques have been implemented in order to enhance the daily workflow in a hospital (Sawaneh et. al., 2018). These digital transformations include improved Wireless Networks, Cloud Solutions, Data Management Systems as well as many other software which their applications may help running the medical devices that are available in several departments of a hospital. Implementing such Information System will lead to the improvements and contribute into a collective system which enables hospitals to work more efficiently such as the Hospital Information System (HIS) (Balaraman, and Kosalram, 2013). Although HIS was primarily used for financial purposes and only partially for patient care

systems, later it became more complex as many hospitals incorporated these specialized systems in several departments (Chatzoglou et. al., 2012) (Mehdipour, and Zerehkafi, 2013) . One of the main departments of a hospital where these digital systems are applied are the radiological departments. They use the data (patient information) which is stored in the HIS, moreover these departments use images created from medical devices such as X-Ray, Computed Tomography, Ultrasound, and Magnetic Resonance Imaging etc. which contain valuable data. This leads to the necessity that a radiological department needs to have its own Information System alongside with the HIS which is internationally known as a Radiology Information System (RIS) (Babić et. al., 2012). RIS is a database designed to support operational workflow and medical analysis within a radiology department. This repository stores patients' personal data and exams carried out, but it has only limited capacity in storing images and dosimetric values which are obtained from

these examinations (Radiologists, 2008). This leads to the issue that images are lost in some cases hence would result in a patient's re-examination and would also mean to expose the patient to higher radiation dose (Palorini et. al., 2014) (Nitrosi et. al., 2014) . Dosimetric information, together with RIS information such as patient data and report outcome could contextualize the dose and consequently help in optimizing the image quality. In addition, dosimetric value information will be stored in the additional database allowing further real patient centered dosimetric value optimization and treatment which enhances clinical accuracy for patients undergoing many ionizing procedures (Palorini et. al., 2014) (Nitrosi et. al., 2014). However, there is another issue to be solved which is, due to the limited capacity of RIS in storing images. To overcome this issue, there are several other software solutions that run alongside the RIS. These techniques will not only solve the limited storage capacity of RIS but instead will avoid unnecessary or unproductive radiation exposure.

One of these techniques is the introduction of the Diagnostic Reference Levels (DRL) which was implemented by the International Commission on Radiological Protection (ICRP) in 1991 (COMMISSION, Radiation protection N° 185, 2018). to ensure adequate doses that a patient may receive in the different diagnoses examinations, a quality management system should be employed (COMMISSION, Radiation protection N° 185, 2018). And as a part of the quality management in many countries DRLs has been established in 1997 and was introduced to the European legislation by the Medical Exposure Directive 97/43/EURATOM (COMMISSION, Radiation protection N° 185, 2018) This was established in order to ensure that doses to which patients are subjected are as low as reasonably possible. However, it is important to mention that DRLs are not dose limits but more of a guide value, mostly utilized to analyze imaging methods which normally result in extremely high patient doses and thus should be adjusted and optimized accordingly (COMMISSION, Radiation protection N° 185, 2018).

Computed Tomography (CT) as one of the useful techniques in diagnostic radiology has a very extensive scope of clinical functions and since due to the CT scan high radiation dose it is a subject of interest for many institutions, therefore internationally it relies heavily on

DRLs (Bosch de Basea et. al., 2015). Since it is generally considered that dose radiation, regardless of the amount, is able to harm the patient, the CT's high radiation dose are a subject of interest for many institutions (Kavanagh et. al. , 2018) (Pearce et. al., 2012).

It has been noticed that hospitals and health institutes in Duhok/KRG - Iraq does not have any DRL standards as well as the mentioned additional repository to store the images and the dosimetric values. Meaning, that Radiologists and Physicians work very hard to manage the examinations without any standards, where they only estimate the patient's dose amount based on their knowledge and experience. Furthermore, in many times they also have to repeat examinations since many patients lose their images and thus have to be re-examined. For these reasons and others, it is not only insufficient and inaccurate; this creates an unnecessary financial burden on the hospitals and it also raises harm for patients since too much radiation can cause severe health risks. Therefore, the aim of this paper was to suggest a DRL model as well as to recommend an additional repository system which provide hospitals in Duhok the necessary platform to manage patient data including images in order to improve the patient's safety as well as helping hospitals and health institutes optimizing the dose radiation management. As a model this paper has taken several CT Scan examples from the Azadi Hospital in Duhok to suggest some possible DRL standards in order to adjust more to the European healthcare enterprise by improving quality and safety.

2. MATERIALS AND METHODS

The CT scan device model Toshiba 64 slice with its software system version "Aquilion V4.61ER004" is currently used in the Azadi Hospital with the company set up configuration. This CT scan system include the standard protocols for different body parts (Head, Neck, Chest, Abdomen, Pelvis, Leg, Chest-to-Pelvis). In this work, all use-cases (samples) were focused primarily on the most used protocols in the Azadi Hospital which included head, chest and abdomen CT examinations. All needed data for this work were self-collected during personal attendance at the CT scan examination sessions. A total number of 70-80 patient data were collected. However, complete CT performance information and examination parameters that

served the purpose of this work, were only obtained from 15 patients as shown in Table 1. In order to process and store these data, an integrated development environment (IDE), such

as Visual Studio or Eclipse is needed. The program and script languages which are recommended are SQL, C# and Python.

Table (1): CT Examination Parameters from Azadi Hospital

Name	Gender	Age	Body Part	Scan Mode	Tube Potential (kV)	Tube Current (mA)	Rotation Time	Total Scan Time	Slice Thickness	CTDI _{vol} (mGy)	DLP (mGy.cm)
Patient 1	F	42	Head	Brain HCT (Helical)	120	300	0.75	18.09	0.5x32	80.5	1503.6
Patient 2	M	70	Head	Brain HCT (Helical)	120	300	0.75	18.44	0.5x32	80.8	1544
Patient 3	M	54	Head	Brain HCT (Helical)	120	300	0.75	17.76	0.5x32	80.3	1499.4
Patient 4	F	49	Head	Brain HCT (Helical)	120	300	0.75	17.73	0.5x32	80	1463.2
Patient 5	M	50	Head	Brain HCT (Helical)	120	300	0.75	17.98	0.5x32	80.8	1491
Patient 6	M	71	Chest	Chest HCT (Helical)	120	268	0.5	19.45	0.5x64	9.9	402.9
Patient 7	F	66	Chest	Chest HCT (Helical)	120	350	0.5	18.77	0.5x64	6.5	292.6
Patient 8	M	54	Chest	Chest HCT (Helical)	120	281	0.5	18.96	0.5x64	8	373.8
Patient 9	M	66	Chest	Chest HCT (Helical)	120	305	0.5	19.12	0.5x64	6.5	292.6
Patient 10	M	73	Chest	Chest HCT (Helical)	120	264	0.5	21.22	0.5x64	10.9	390.7
Patient 11	F	60	Abdomen	Abdomen-C HCT (Helical)	120	460	0.5	19.55	0.5x64	35	1581.2
Patient 12	F	29	Abdomen	Abdomen-C HCT (Helical)	120	403	0.5	18.98	0.5x64	30.6	1020.3
Patient 13	F	34	Abdomen	Abdomen-C HCT (Helical)	120	500	0.5	19.61	0.5x64	38	2036.1
Patient 14	M	71	Abdomen	Abdomen-C HCT (Helical)	120	328	0.5	18.71	0.5x64	24.2	1646.1
Patient 15	F	56	Abdomen	Abdomen-C HCT (Helical)	120	425	0.5	19.42	0.5x64	32.5	1245

To start, the patients had to give his/her personal information to the HIS which contained name, age, gender, weight, etc. The HIS then assigned a Patient ID that will function as the unique identifier. After that the patient normally

is send according to the diagnosis to a medical device (e.g. CT scan) which created an image of the correspondent body part as shown in the diagram of Figure 1.

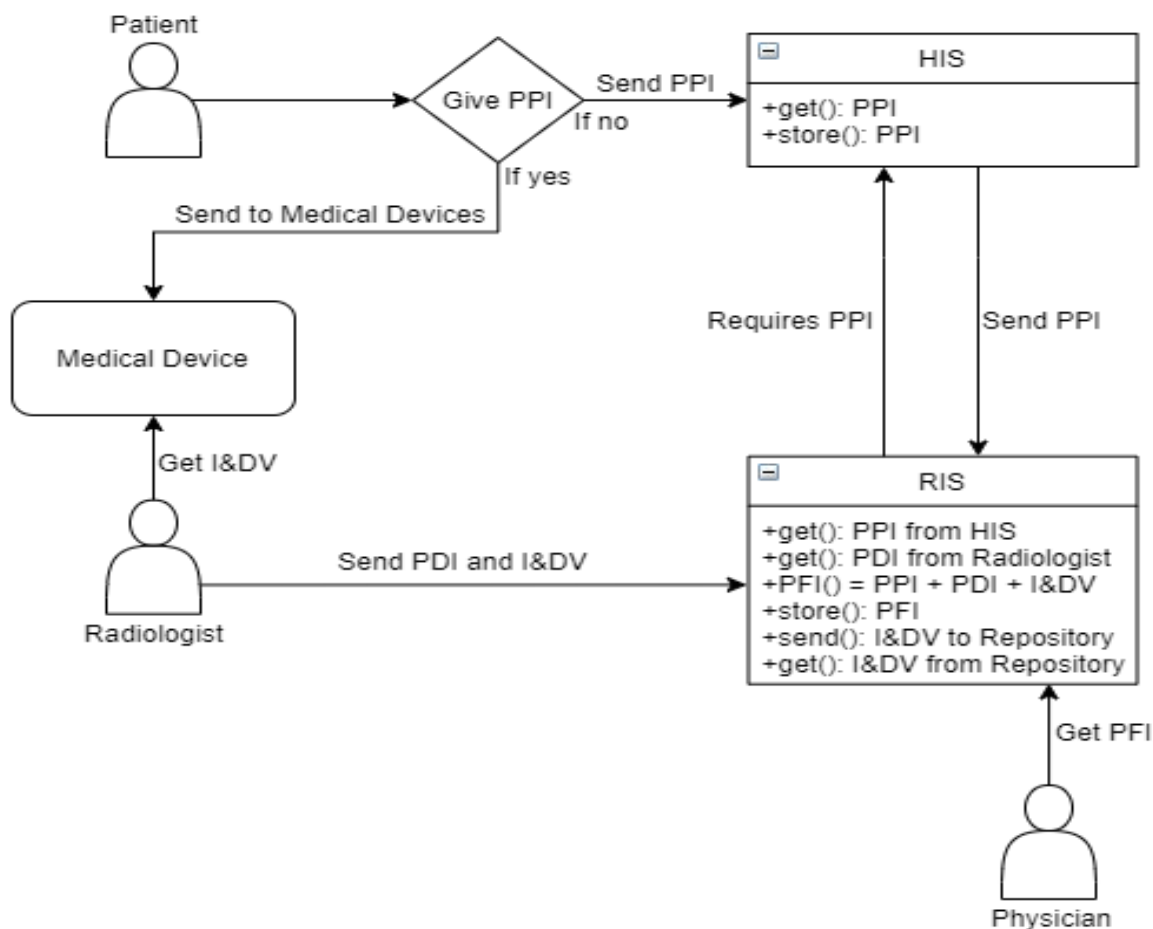


Fig. (1): Flow Chart Diagram developed to display a typical use-case scenario in the hospital

HIS = Hospital Information System / RIS = Radiology Information System / PPI = Patient Personal Information / PDI = Patient Detailed Information / PFI = Patient Full Information / I = Imagery / DV = Dosimetric Values / Medical Devices can include: X-Ray, Computed Tomography, Magnetic Resonance Imaging, Ultrasound etc.

In addition, the multi-detector CT scanner recorded several data, such as the dosimetric values. The three main dose indices are Computed Tomography Dose Index volume ($CTDI_{vol}$) measured in milligrays (mGy), the Dose Length Product (DLP) measured in milligrays per centimeter (mGy-cm) and the Effective Dose (ED) measured in millisieverts (mSv) as was shown in Table 1 (Granata et. al., 2014).

However, storing these values together with the imagery was proven to be an issue since the available storage space was not enough to contain all these documents, which lead to the fact that patients had to carry their x-rays with

them each time they had to revisit the hospital, thus an additional repository was needed in order to fully store all relevant patient documents.

As mentioned above the CT scanner has been configured by the suppliers and it seemed that it has been limited due to the company's policy in addition, to unavailability of necessary software and thus, a number of dosimetric values were not shown. Therefore, to determine all the values needed in order to set a possible DRL standard as well as revealing the difference in the results between patients using the same body part, a number of calculations were carried out. One of these values to be calculated was the estimated length (L) measured in centimeter (cm) which was based on the $CTDI_{vol}$ and DLP generated by the CT scanner according to the following equation (Ekpo et. al., 2018) (Saravanakumar et. al., 2014).

$$DLP = CTDI_{vol} \times L \quad (1)$$

The second value to be calculated was the Effective Dose (ED) which represents the equivalent whole-body dose according to the

following equation (Deak et. al., 2010) (Ekpo et. al., 2018)

$$ED = DLP \times k \quad (2)$$

Where k is the tissue weighting factor based on the scanned body region. Conversion factors

for adults of various ages are shown in Table 2 as described and recommended in the ICRP publication 60 & 103 (Eckerman et. al., 2012) (VALENTIN, 2007) (Kobayashi et. al., 2019).

Table (2): Conversion Factors for adults

Scanned body parts	k (tissue weighting factor)
Head / Neck	0.0031
Head	0.0021
Neck	0.0059
Chest	0.014
Abdomen	0.015
Trunk	0.015

The values of $CTDI_{vol}$, DLP, L and ED for each examination sample were calculated and recorded. These value calculations and dosimetry technique was based on the methods proposed by European Guidelines and also according to the conditions and recommendations of the ICRP (VALENTIN, 2007) (COMMISSION, 2000) (Saravanakumar et. al., 2014). Additional values that were calculated in this work, was to determine the percentiles of the three dosimetric values. It is important to mention that image quality improves proportional with the increasing percentage of dose radiation and vice versa. Therefore, the most commonly used percentile as an indicator in Europe to do the DRL calculations for each examination are 25th, 50th and 75th percentile (Ekpo et. al., 2018) (Nakada Y et. al., 2018) (Saravanakumar et. al., 2014)

While the 25th percentile values are normally used to assess the lowest dose levels for the price of significantly deteriorated image quality, the 50th percentile on the other hand, functions as median and provides dose levels that institutes should strive towards to. However, the 75th percentile values are also the most effective

when establishing the DRL function also as proposed dose values which should not be exceeded (VALENTIN, 2007). Most European countries use the 75th percentile values thus this work will estimate the DRL values using the same percentile as the European countries do in order to be able to compare them with each other (Health, 2018) (Richard Veit, and Burkhard Bauer, 2019).

Development of a DRL model

In this work, the 75th percentile of the dosimetric values $CTDI_{vol}$, DLP and ED were then used to calculate the DRL values. Then, these new calculated DRL values were compared with the ones of a number of European countries to identify whether the obtained data exceeded their limits and thus to suggest the needs for adjustment. Finally, a further developed flow chart was designed to integrate an Information System model.

3. RESULTS AND DISCUSSION

The results of the first step which involved the calculation of length L, are summarized in Table 3. These results were obtained by extracting the dosimetric values from the additional database via the RIS.

Table (3): Dosimetric Values and Patient's Length

Name	CTDI _{vol} (mGy)	DLP (mGy.cm)	Length (cm)
Patient 1	80.5	1503.6	18.68
Patient 2	80.8	1544	19.11
Patient 3	80.3	1499.4	18.67
Patient 4	80	1463.2	18.29
Patient 5	80.8	1491	18.45
Patient 6	9.9	402.9	40.70
Patient 7	6.5	292.6	45.02
Patient 8	8	373.8	46.73
Patient 9	6.5	292.6	45.02
Patient 10	10.9	390.7	35.84
Patient 11	35	1581.2	45.18
Patient 12	30.6	1020.3	33.34
Patient 13	38	2036.1	53.58
Patient 14	24.2	1646.1	68.02
Patient 15	32.5	1245	38.31

From this Table it can be noticed that L varies within the same protocol, where the patient samples from 1 to 5 represented the Head protocol, the patient samples from 6 to 10 represented the Chest protocol and the patient samples 11 to 15 represented the Abdomen protocol. For example, the two main dosimetric values for Patient 1 were obtained by the CT scanner with 80.5 mGy and 1503.6 mGy.cm respectively. Then, by applying equation (1), the estimated L was indicated 18.68 cm and all the other samples followed the same procedures.

The shown result in the above Table indicated, that the observed amount of the DLP

value increased proportional with the raising length L, which led to the fact that the amount of the DLP could be controlled and limited by the radiologist/physician depending on the length of the body part that was selected and estimated (BUSHBERG et. al., 2012).

The results of the following step represented the Effective Dose (ED) measured by applying equation (2) with the use of the k factor for the Head was 0.0021, for the Chest was 0.014 and for the Abdomen was 0.015 where all results are represented in Table 4.

Table (4): The Effective Dose for each Patient

Name	DLP (mGy.cm)	K Factor	ED (mSv)
Patient 1	1503.6	0.0021	3.16
Patient 2	1544	0.0021	3.24
Patient 3	1499.4	0.0021	3.15
Patient 4	1463.2	0.0021	3.07
Patient 5	1491	0.0021	3.13
Patient 6	402.9	0.014	5.64
Patient 7	292.6	0.014	4.10
Patient 8	373.8	0.014	5.23
Patient 9	292.6	0.014	4.10
Patient 10	390.7	0.014	5.47
Patient 11	1581.2	0.015	23.72
Patient 12	1020.3	0.015	15.30
Patient 13	2036.1	0.015	30.54
Patient 14	1646.1	0.015	24.69
Patient 15	1245	0.015	18.68

The results revealed that the ED amount also increased proportional to the amount of the DLP, which described the dose amount that each patient received during the scanning (Ekpo et. al., 2018) (Huda et. al., 2008).

According to the results shown in Table 3 and 4, it was possible to consider that the selected length played the main role in the dose

amount that was given to any patient. From these results it appeared that both, the length and the ED in most cases are higher in comparison to other countries, meaning that a higher length correlated to a higher dose amount thus exposing patients to a greater harm of radiation.

The result of different samples showed how the used method could help identify where

protocols are probably not standardized and to suggest where further optimization actions should be taken. The factors that affect and help to establish the DRL are $CTDI_{vol}$, DLP and ED (Ekpo et. al., 2018) (Saravanakumar et. al., 2014).

Actually, all three factors are routinely used for comparing exposures from different scanning protocols, for setting diagnostic reference levels (DRLs), and for dose optimization. DRLs allow radiology departments to compare their dose levels with regional or national standards. These levels are expected not to be exceeded for

standard procedures when good and normal practice regarding diagnostic and technical performance is applied (Granata et. al., 2014).

Development of a DRL Model

In order to design a DRL model in our hospitals, the dosimetric values were used to determine the commonly used percentiles (25th, 50th and 75th) applied by the European countries for all three protocols as shown in Table 5. The importance of providing these data, is for any further studies that wishes to develop a DRL model using any of these percentiles.

Table (5): Percentiles of Dosimetric Values

Body region	CTDI _{vol} (mGy) Percentile values			DLP (mGy.cm) Percentile values			Effective Dose (mSv) Percentile values		
	25 th	50 th	75 th	25 th	50 th	75 th	25 th	50 th	75 th
Head	80.3	80.5	80.8	1491	1499	1504	3.13	3.15	3.16
Chest	6.5	8	9.9	292.6	373.8	390.7	4.1	5.23	5.47
Abdomen	30.6	32.5	35	1245	1581	1646	18.68	23.72	24.69

The DRL standards were then calculated by using the most frequently used percentiles 75th of the dosimetric values, a value that is used by most of the European countries. These calculations were made according to the European Guidelines and also according to the conditions and recommendations of the ICRP (VALENTIN, 2007) (COMMISSION, 2000) (Saravanakumar et. al., 2014).

As mentioned before, the selected length could be considered as the main role in the dose amount that was given to any patient. Therefore, the European countries set an average length for each specific part of the body region. Thus, in this study it was recommendable to do so as well. Using the 75th percentile, the length values for the samples have been calculated by measuring the ratio of the two dosimetric values DLP and $CTDI_{vol}$ as shown in Table 6.

Table (6): Length values using 75th percentile

Body region	CTDI _{vol} (mGy)	DLP (mGy.cm)	L (DLP/CTDI)(cm)
Head	80.8	1503.6	18.61
Chest	9.9	390.7	39.46
Abdomen	35	1646.1	47.03

The results of Table 6 made clear that the length of the Head with 18.61 cm, Chest with 39.46 cm and Abdomen with 47.03 cm were exceeding the ones from other European countries. For example, in Germany as one of the leading countries in this domain the average length for the Head, Chest and Abdomen region is almost 14.17, 35 and 24 cm respectively (Stamm et. al., 2017).

Most European countries use the 75th percentile values due to the fact that DRLs

should be made accessible for common studies with commensurate dose values. Therefore, collaborations between authorities and professional associations have been established with the purpose of creating national DRLs by using the 75th percentile of the examined distribution of patient doses in their countries (Health, 2018) (Richard Veit, and Burkhard Bauer, 2019).

It is important to mention that these DRL values cannot be considered as standard values

unless they are in a suitable range of other reported DRL ranges. Therefore, these study values were further compared with other

standardized DRL values from several selected European countries as shown in Table 7.

Table (7): Dosimetric values of selected European countries

Body region	Head			Chest			Abdomen		
	CTDI _{vol}	DLP	ED	CTDI _{vol}	DLP	ED	CTDI _{vol}	DLP	ED
This study	81	1504	3.16	10	391	5.47	35	1646	24.69
Germany	60	850	1.79	10	350	4.90	15	360	5.40
Sweden	60	1000	2.10	9	350	4.90	11	550	8.25
Italy	69	1312	2.76	15	569	7.97	18	550	8.25
Slovenia	56	865	1.82	13	420	5.88	16	550	8.25
Czech Republic	65	1100	2.31	15	500	7.00	19	750	11.25
Belgium	50	900	1.89	8	260	3.64	10	490	7.35
France	58	1050	2.21	20	500	7.00	25	650	9.75
Bulgaria	60	1000	2.10	16	500	7.00	18	470	7.05
Poland	60	1050	2.21	30	650	9.10	35	780	11.70
Finland	55	800	1.68	9	290	4.06	12	560	8.40

The values in Table 7 display the CTDI_{vol}, DLP and ED range domain for Head, Chest and Abdomen. The range domain was determined from the lowest to the highest value for all selected European countries. This was valid for CTDI_{vol}, DLP and ED. For example, in the Head protocol, the CTDI_{vol} value obtained in this study was 81 while the lowest CTDI_{vol} value for the same protocol reported in Belgium was 50 and the highest value reported in Italy was 69. This made clear that the determined value of this study of 81 was exceeding the range domain compared to the selected European countries. The DLP value for the Head protocol obtained in this study was 1504. This value also exceeded the range domain in the selected countries where the lowest DLP value in Finland was 800 and the highest value in Italy was 1312. As for the ED,

the value of the Head protocol of this study was 3.16 while the lowest value reported in Finland was 1.68 and the highest value reported in Italy was 2.76. As a result, the correspondent DRL values of this study were also high due the three dosimetric values. However, it is interesting to mention that the result of the Chest area showed that it is well within the middle of the range domain of the selected countries. The results of the dosimetric values for Abdomen from this study were also very high compared to the ones from the other countries. Thus, standard limitations are needed in order to be able to reduce the patient's dose amount. A more detailed range display from the lowest to the highest values of CTDI_{vol}, DLP and ED within all three protocols Head, Chest and Abdomen is shown in Figure 2.

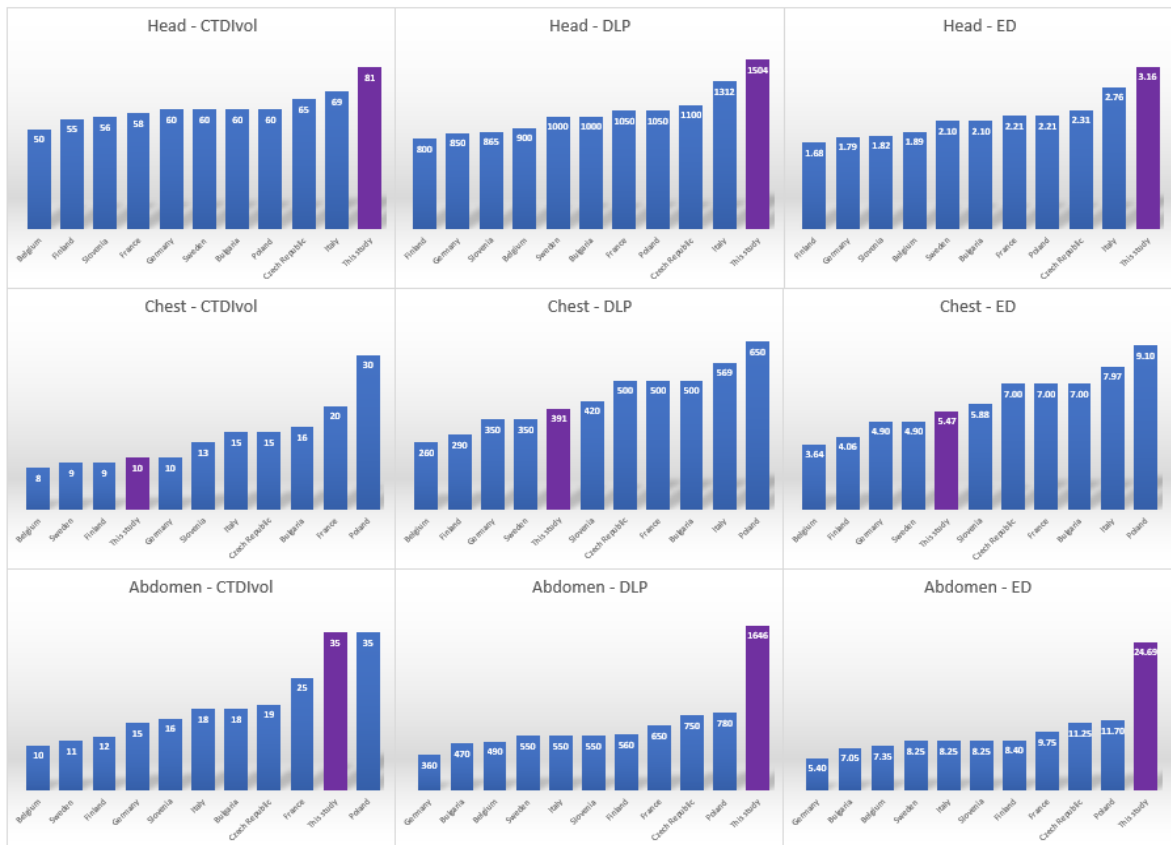


Fig. (2): Dosimetric value range domain for Head, Chest and Abdomen

The results of the Length obtained in this study for the three protocols were also compared to the selected European countries as displayed in Table 8.

Table (8): Length Comparison

Body region	Head	Chest	Abdomen
Dosimetriv values	L (DLP/CTDI)	L (DLP/CTDI)	L (DLP/CTDI)
This study	18.57	39.10	47.03
Germany	14.17	35.00	24.00
Sweden	16.67	38.89	50.00
Italy	19.01	37.93	30.56
Slovenia	15.45	32.31	34.38
Czech Republic	16.92	33.33	39.47
Belgium	18.00	32.50	49.00
France	18.10	25.00	26.00
Bulgaria	16.67	31.25	26.11
Poland	17.50	21.67	22.29
Finland	14.55	32.22	46.67

A more detailed range display from the lowest to the highest Length within all three protocols Head, Chest and Abdomen is shown in Figure 3.

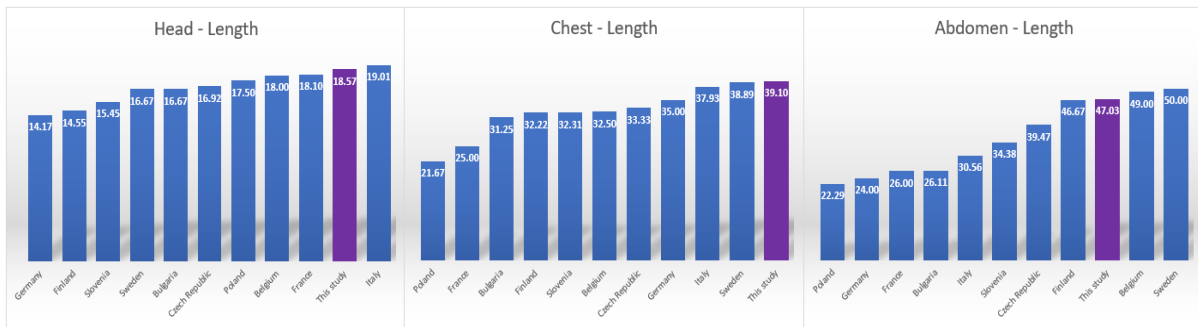


Fig. (3): Detailed Length Display

Analyzing this figure, the following notes may generally be extracted. For example, while the dosimetric values for Chest are within average range domain, the Length is only a little higher compared with the other countries and therefore could be reduced. The real problematic case was reflected in the Head and Abdomen protocol since the dosimetric values and the

Length were very high. A dose amount should be adjusted, by implementing a suggested DRL.

With help of this new designed DRL model, a new control mechanism model was constructed which functioned as an interface between the RIS and the Radiologist/Physician as shown in Figure 4.

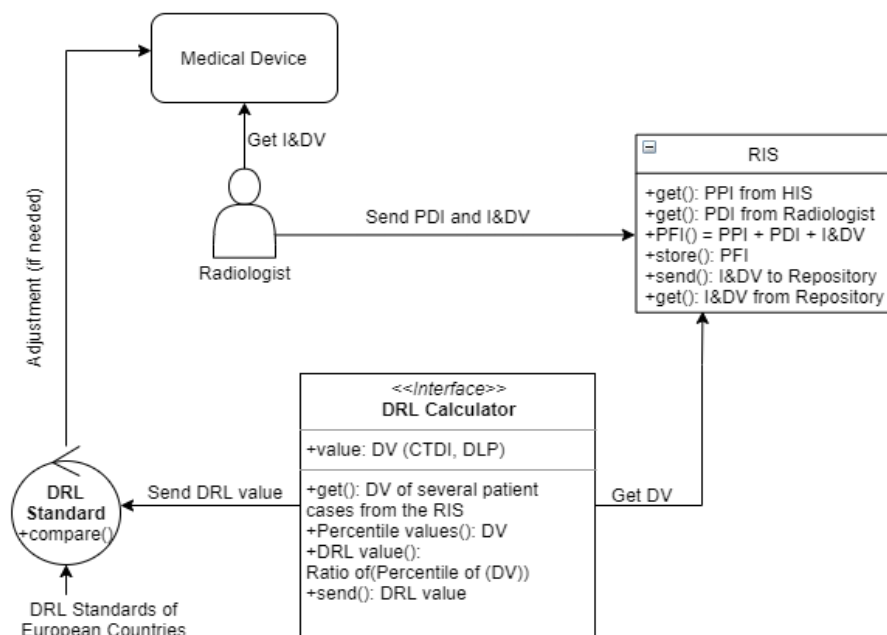


Fig. (4): Suggested DRL Control Model

Implementing this model, the interface will send the DRL value to the control unit “DRL Standard” which then evaluated it by comparing them with the standard values of the European countries. If the DRL value exceeded the European standard values, then the Radiologist/Physician is asked by the system, more precisely by the software which runs the CT scanner, to adjust the estimated length, since

the results made clear that the selection of a body region’s length has a direct impact of the dose amount. However, if the calculated DRL value was within the European standard values, then the radiologist/physician could continue with the process.

A final Flow Chart as shown in Figure 5 was designed, which may run the CT scan procedure effectively as well as control and suggest adjustments

of the dose amount. It is worth mentioning that an extra repository system was added to the model that

should help in solving the short storage capacity of the RIS as mentioned previously.

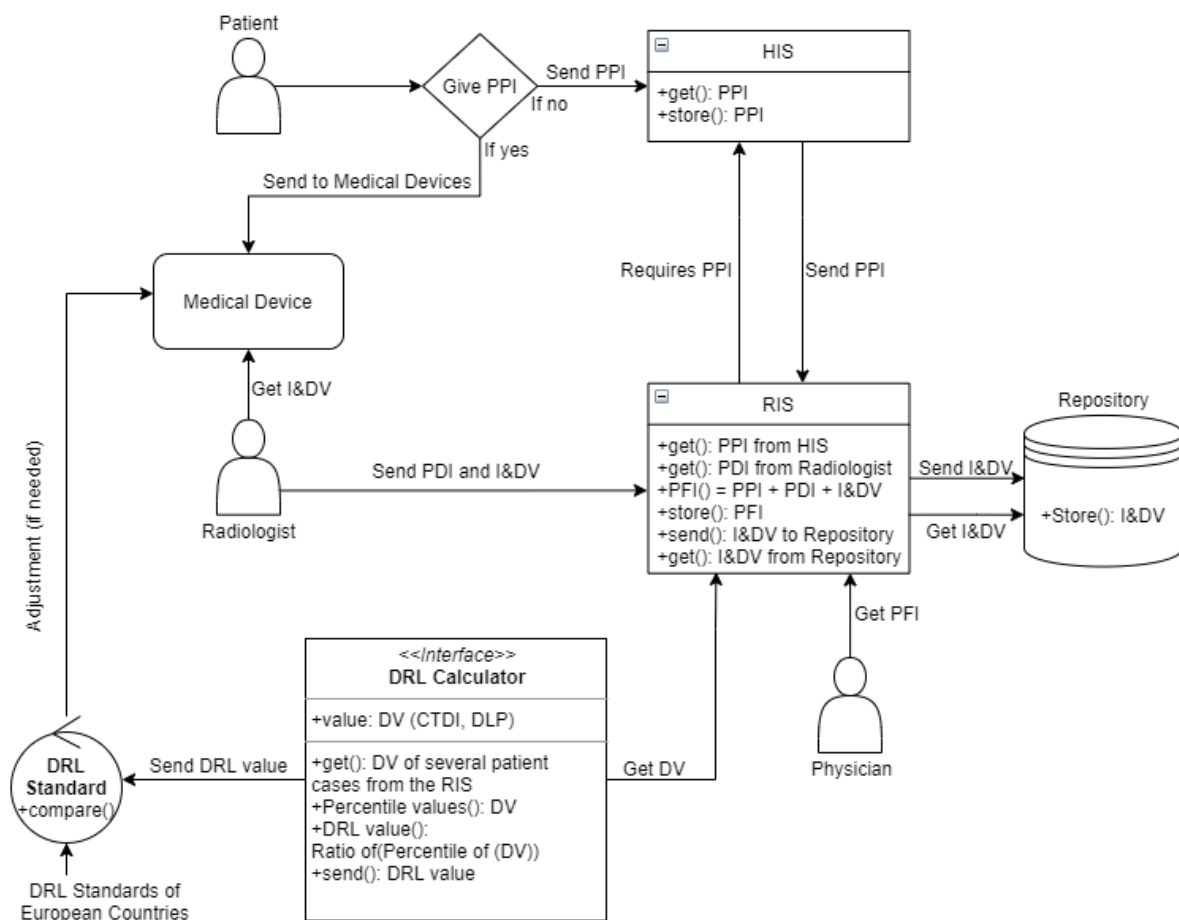


Fig. (5): Enhanced Flow Chart Diagram of a typical use-case scenario in the hospital

To this end, integrating the new Information System Model will improve the benefit of such important medical equipment and on the same time minimizing the effects of harmful and unnecessary doses of radiation that a patient may be exposed to while it may also prevent unnecessary financial expenditure for the hospital.

4. CONCLUSION

This work involved the integration of an Information System for suggesting a DRL model that may ensure the safety of a patient by implementing two solution methods. The first was to set an additional repository to overcome the capacity limitations of the hospital’s data storage. The second solution was to create a safe radiation management system through

suggesting a DRL model at the CT scan unit in Azadi hospital.

The results revealed that the dosimetric values were much higher than those internationally reported. Therefore, from these findings it may be concluded that such an Information System is of a necessity for patient’s safety as well as for supporting the Radiologists/Physicians to perform radiological examinations in a safer manner in addition; to reducing financial costs and improvement of the hospital daily workflow.

5. ACKNOWLEDGEMENT

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