RESEARCH ARTICLE

MEDICAL PHYSICS

Patient-specific fetal radiation dosimetry for pregnant patients undergoing abdominal and pelvic CT imaging

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Funding information

National Natural Science Foundation of China, Grant/Award Number: 32227801; Swiss National Science Foundation, Grant/Award Number: SNSF 320030_176052

Abstract

Background: Accurate estimation of fetal radiation dose is crucial for riskbenefit analysis of radiological imaging, while the radiation dosimetry studies based on individual pregnant patient are highly desired.

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Purpose: To use Monte Carlo calculations for estimation of fetal radiation dose from abdominal and pelvic computed tomography (CT) examinations for a population of patients with a range of variations in patients' anatomy, abdominal circumference, gestational age (GA), fetal depth (FD), and fetal development.

Methods: Forty-four patient-specific pregnant female models were constructed based on CT imaging data of pregnant patients, with gestational ages ranging from 8 to 35 weeks. The simulation of abdominal and pelvic helical CT examinations was performed on three validated commercial scanner systems to calculate organ-level fetal radiation dose.

Results: The absorbed radiation dose to the fetus ranged between 0.97 and 2.24 mGy, with an average of 1.63 ± 0.33 mGy. The CTDI_{vol}-normalized fetal dose ranged between 0.56 and 1.30, with an average of 0.94 ± 0.25 . The normalized fetal organ dose showed significant correlations with gestational age, maternal abdominal circumference (MAC), and fetal depth. The use of ATCM technique increased the fetal radiation dose in some patients.

Conclusion: A technique enabling the calculation of organ-level radiation dose to the fetus was developed from models of actual anatomy representing a range of gestational age, maternal size, and fetal position. The developed maternal and fetal models provide a basis for reliable and accurate radiation dose estimation to fetal organs.

KEYWORDS

computational model, CT, fetal radiation dose, Monte Carlo, pregnant patients

1 | INTRODUCTION

Pregnant females might be exposed to ionizing radiation during radiological procedures performed for clinical diagnosis of acute abdominal pain, trauma cancer and a number of other clinical indications. As the embryo/fetus presents significantly higher radiosensitivity than adults,¹ the International Commission on Radiological Protection (ICRP) reported that radiation doses above 100 mGy may lead to deterministic

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effects to the fetus, such as malformation, death, and mental retardation.² In addition, the ICRP publication 103 suggested that fetal radiation doses lower than 100 mGy cannot be used as an argument to stop pregnancy, a decision that should be carefully studied from different perspectives.³ It is important to estimate fetal organ doses from diagnostic radiological examinations to understand the risks to the developing fetus from ionizing radiation. Due to ethical concerns, the fetal radiation dose can hardly be directly measured and has become a challenge for medical physicists. MEDICAL PHYSICS

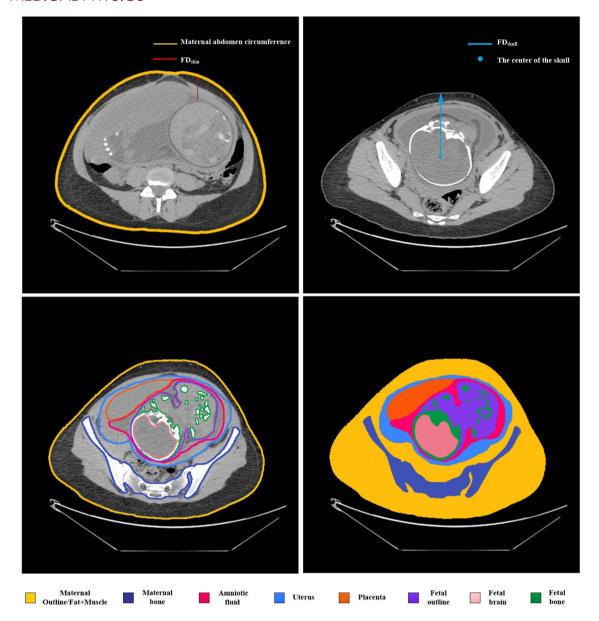


FIGURE 1 CT images of a pregnant female patient at 35 weeks of gestation showing the segmented organs and measurements of maternal abdominal circumference, fetal skin depth (FD_{skin}), and fetal skull depth (FD_{skin}) distances.

A number of studies have reported on the use of Monte Carlo (MC) simulations to estimate fetal radiation dose,^{4–10} where fetal dosimetry was performed using a reference model or a maternal-size-adjusted reference model without including the individual anatomical characteristics of the fetus, hence introducing an extra error in radiation dosimetry.⁴ For example, Maynard et al.¹¹ constructed eight reference models for pregnant women. The ICRP Publication 145¹² also published a series of mesh-type reference computational phantoms. With the increasing interest in personalized radiation dosimetry among the scientific community, the construction of personalized computational models has become popular.^{13,14} Angel et al.¹⁵ constructed patient-

specific voxel models for 24 pregnant patients for fetal dose calculation in abdominal CT examinations, but the fetal organ-level radiation dose was immeasurable because of the limited number of identified fetal organs. Since the radiosensitivity and radiation risks of developing fetal organs vary across the different gestational ages, organ-scale radiation dosimetry for the fetus is crucial and highly desired for epidemiological studies aiming to correlate between conceptus radiation exposure and organ-specific childhood cancer after birth. In addition, most of the aforementioned previous work was performed using fixed tube current (FTC) acquisition protocols. As automatic tube current-modulated (ATCM) CT scanning becomes popular, it is important to

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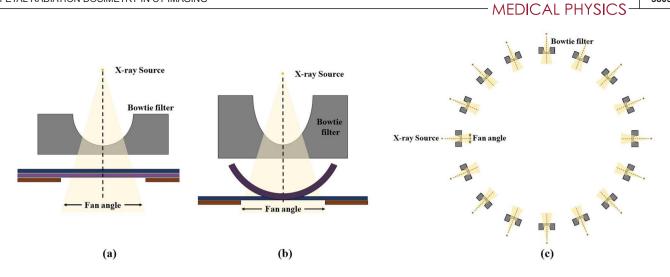


FIGURE 2 Geometry of three validated commercial CT scanner models: (a) GE Discovery CT 750 HD scanner, (b) Siemens SOMATOM Definition Edge scanner, and (c) United Imaging Healthcare uCT 760 128-slice CT scanner.

estimate fetal dose from different scanning protocols across a variety of CT scanners.

In this work, we constructed a series of pregnant female models at different gestational ages based on CT images of real patients and performed MC simulations to calculate the radiation dose delivered to the fetus. Three previously validated CT scanner systems, including the GE Discovery CT 750 HD, the Siemens SOMATOM Definition Edge, and the United Imaging Healthcare uCT 760 128-slice CT, were incorporated into the MC simulations. The fetal radiation dose from abdominal/pelvic CT examinations on five scanners, including the Siemens SOMATOM Force, the Philips Brilliance 40, the Siemens Definition AS, the GE LightSpeed VCT, and the GE Discovery CT750 HD, were calculated and compared. The correlations between fetal organ dose from CT examinations and various attributes such as scanning protocols, individual anatomical characteristics, gestational age, intra-uterine position, and fetal size were investigated.

2 | MATERIALS AND METHODS

2.1 | Patient-specific computational models

A total of 44 patient-specific computational models were constructed based on clinical CT images of pregnant patients acquired at Geneva University Hospital (HUG), Switzerland. These patients were initially admitted to the emergency department of HUG because of acute abdominal pain and underwent ultrasound examinations upon admission, which however did not yield an etiology or conclusive diagnosis. MRI examination was not possible either owing to lack of slots or patients presenting very late in the evening. Hence, the patients underwent abdominal and pelvic low-dose CT examinations. CT images of pregnant patients were segmented using Photoshop software (San Jose, California, USA), while the NURBS-based organ surfaces were reconstructed using Rhinoceros software (Seattle, Washington, USA). The construction of patient-specific computational models for pregnant females based on clinical CT images has been described in detail in the prior article.¹⁶ This prior article dealt with development of digital fetus library and the measurement of anatomical parameters, whereas in this manuscript, we report on the radiation dosimetry study for fetus and pregnant patient in abdominal and pelvic CT examinations. The measured anatomical parameters include femoral length (FL), humerus length (HL), biparietal diameter (BPD), head circumference (HC), abdominal circumference (FAC), fetal skin depth (FD_{skin}), and fetal skull depth (FD_{skull}). The constructed abdominal models of pregnant female subjects were named Maternal_Abdominal_Model-X (MAM-X) where X refers to the model ID. Figure 1 illustrates the manual methods of image segmentation and measurements of the maternal abdominal circumference and intra-uterine position of FD_{skin} and FD_{skull}. MAC was taken as the circumference of the pregnant woman's belly button, while FD_{skin} and FD_{skull} were calculated as the shortest distances between the most front of the fetal skin and the 3D center of the fetal skull to the skin surface of the pregnant woman, respectively. All above-mentioned feature points and data were measured using the Rhinoceros software.

2.2 | Monte Carlo simulation and radiation dose calculations

The study included three clinical CT scanner systems, namely, a GE Discovery CT 750 HD scanner,^{4,17} a

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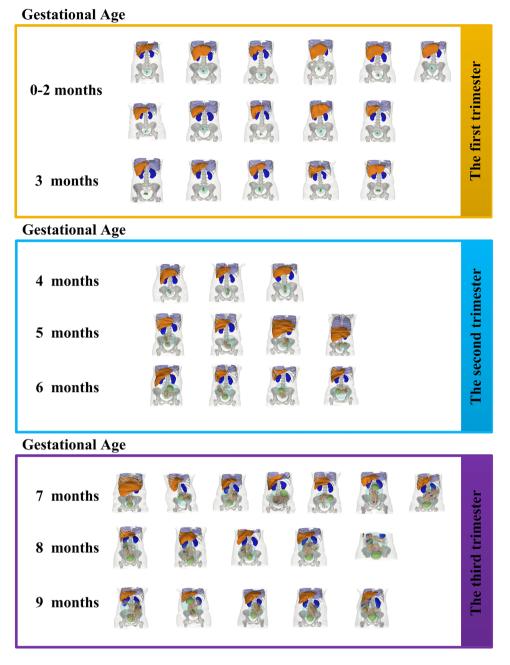


FIGURE 3 3D visualization of the developed patient-specific computational phantoms at different gestational ages/trimesters. The outer outline of pregnant patients and the uterus are transparent for better visualization of internal organs.

Siemens SOMATOM Definition Edge scanner,¹⁸ and a the United Imaging Healthcare uCT 760 128-slice CT scanner,¹⁹ which had been previously incorporated into the MCNPX Monte Carlo code²⁰ for calculations of radiation dose of various organs, however, not for pregnant patients. The geometries of the three CT gantries, including the source to detector distance, focal spot size, fan angle, etc., had been taken into account in the simulation of corresponding CT scanners as shown in Figure 2. Simulations of x-ray photon transport were performed to yield dose calculations based

on 44 pregnant-female phantoms where the fetal radiation dose refers to the total deposited energy in the fetal body divided by the body weight of the fetus. For each phantom, the simulated energy deposition in 25 fetal organs and maternal body were recorded and used for the radiation dosimetry calculations. For each patient phantom and for each organ, the simulated radiation doses were averaged across all three simulated CT scanners. The detailed data processing and correlation analysis methods are provided in Supplemental Materials.
 TABLE 1
 Anatomical measurements for 44 pregnant patients and their fetuses.

Model Number	GA (weeks)	MAC (cm)	CT scanners	Mean CTDI _{vol} (mGy)	Mean effective mAs	Scan length (cm)	FL (mm)	HL (mm)	BPD (mm)	HC (mm)	FAC (mm)	FD _{skull} (mm)	FD _{skin} (mm)
MAM-1	8	84.63	GE Discovery CT750 HD	1.78	22.88	42.5	_	_	_	_	_	_	-
MAM-2	8	79.35	GE Discovery CT750 HD	1.78	22.88	41.5	_	-	-	-	-	-	-
MAM-3	8	76.90	GE Discovery CT750 HD	1.78	22.88	41.9	-	-	-	-	-	-	-
MAM-4	8	70.40	GE Discovery CT750 HD	1.78	22.88	45.5	-	-	-	-	-	-	-
MAM-5	8	78.98	Philips Brilliance 40	9.59	162.51	47.3	-	-	-	-	-	-	-
MAM-6	8	94.09	GE Discovery CT750 HD	1.78	22.88	47.5	-	-	-	-	-	-	-
MAM-7	8	98.46	GE Discovery CT750 HD	1.78	22.88	44.5	_	-	-	-	-	-	-
MAM-8	8	80.99	GE Discovery CT750 HD	1.78	22.88	41.5	_	-	-	-	-	-	-
MAM-9	8	93.33	GE Discovery CT750 HD	1.78	22.88	49.9	-	-	-	-	-	-	-
MAM-10	8	86.13	GE Discovery CT750 HD	1.78	22.88	41.5	-	-	-	-	-	-	-
MAM-11	8	117.80	GE Discovery CT750 HD	1.78	22.88	43.9	_	-	-	_	_	-	-
MAM-12	10	119.00	GE Discovery CT750 HD	1.78	22.88	50.5	-	-	8.69	33.5	_	-	-
MAM-13	10	77.58	GE Discovery CT750 HD	1.78	22.88	43.5	-	-	8.87	32.8	-	-	-
MAM-14	10	93.25	GE Discovery CT750 HD	1.78	22.88	43.9	-	-	9.09	33.0	-	-	-
MAM-15	12	98.99	GE LightSpeed VCT	1.81	22.90	47.9	-	-	9.11	32.7	-	-	-
MAM-16	12	89.18	SIEMENS Definition AS	10.82	141.93	43.9	-	-	9.15	33.4	-	-	-
MAM-17	15	89.28	GE Discovery CT750 HD	1.78	22.88	39.5	30.76	29.13	37.95	134.1	-	-	-
MAM-18	15	122.30	SIEMENS SOMATOM Force	2.94	43.03	48.7	30.67	29.13	36.97	132.9	-	-	-
MAM-19	16	108.30	GE Discovery CT750 HD	1.78	22.88	45.5	32.88	31.31	37.34	133.5	-	-	-
MAM-20	17	92.04	GE Discovery CT750 HD	1.78	22.88	45.9	32.58	31.37	46.91	171.0	151.6	90.5	41.87
MAM-21	18	82.09	GE Discovery CT750 HD	1.78	22.88	43.9	30.97	29.91	48.64	162.8	148.9	85.0	51.42
MAM-22	20	111.00	GE Discovery CT750 HD	1.78	22.88	64.7	34.93	33.33	53.25	175.9	202.6	119.7	63.71
MAM-23	20	112.60	GE Discovery CT750 HD	1.78	22.88	47.9	34.23	34.75	51.65	193.1	189.3	116.0	68.81
MAM-24	21	89.33	GE Discovery CT750 HD	1.78	22.88	41.3	36.04	34.62	52.15	184.4	179.2	82.9	38.36
MAM-25	21	87.93	GE Discovery CT750 HD	1.78	22.88	37.9	31.92	32.94	52.70	188.1	158.0	88.7	58.28

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TABLE 1 (Continued)

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Model Number	GA (weeks)	MAC (cm)	CT scanners	Mean CTDI _{vol} (mGy)	Mean effective mAs	Scan length (cm)	FL (mm)	HL (mm)	BPD (mm)	HC (mm)	FAC (mm)	FD _{skull} (mm)	FD _{skin} (mm)
MAM-26	22	101.90	GE Discovery CT750 HD	1.78	22.88	46.3	42.87	39.84	64.51	218.8	202.1	92.2	54.89
MAM-28	23	98.08	SIEMENS SOMATOM Force	1.51	22.17	47.9	46.64	43.25	68.00	233.2	229.6	96.8	40.11
MAM-29	25	109.50	GE Discovery CT750 HD	1.78	22.88	46.5	68.97	64.31	94.13	321.8	349.2	106.5	24.59
MAM-30	25	92.21	GE Discovery CT750 HD	1.78	22.88	46.5	45.82	39.48	69.19	230.9	233.9	73.1	30.09
MAM-31	25	96.01	GE LightSpeed VCT	1.81	22.90	46.9	58.82	54.92	83.68	300.7	277.6	77.6	37.59
MAM-32	25	116.10	GE Discovery CT750 HD	1.78	22.88	47.5	46.39	43.31	63.81	232.9	242.6	173.9	61.45
MAM-33	26	111.60	GE Discovery CT750 HD	1.78	22.88	44.9	57.28	52.49	75.75	261.5	248.3	101.9	47.62
MAM-34	26	114.00	GE LightSpeed VCT	1.81	22.90	49.9	49.51	40.24	66.86	245.1	211.4	132.0	84.73
MAM-35	28	88.92	GE Discovery CT750 HD	1.78	22.88	43.9	55.96	51.31	76.38	263.5	242.8	63.2	19.80
MAM-36	29	127.90	GE LightSpeed VCT	1.81	22.90	48.1	62.93	58.70	83.24	278.7	264.1	149.5	78.27
MAM-37	29	89.20	GE Discovery CT750 HD	1.78	22.88	47.5	56.15	48.33	78.33	289.3	256.7	71.2	22.89
MAM-39	30	97.68	GE Discovery CT750 HD	1.78	22.88	42.9	61.75	51.21	82.56	285.6	257.4	92.9	22.71
MAM-41	32	114.10	GE Discovery CT750 HD	1.78	22.88	48.5	64.51	61.12	88.30	309.3	278.3	124.0	40.84
MAM-42	33	118.20	GE Discovery CT750 HD	1.78	22.88	46.9	60.91	55.62	85.05	296.2	244.5	117.4	46.23
MAM-43	33	124.50	GE Discovery CT750 HD	1.78	22.88	48.3	61.20	55.62	85.65	295.3	239.8	158.9	52.17
MAM-44	35	106.10	GE Discovery CT750 HD	1.78	22.88	42.5	72.68	56.65	88.53	321.9	332.6	143.5	46.92
MAM-45	35	108.60	GE Discovery CT750 HD	1.78	22.88	47.9	65.95	55.79	84.75	311.7	309.2	116.3	44.88
MAM-46	35	98.80	GE LightSpeed VCT	1.81	22.90	48.3	67.54	58.31	97.02	341.5	327.7	104.6	35.1

2.3 | Fetal dose from automatic tube current modulation (ATCM)

With the advancements of ATCM on modern CT scanners, protocols using this technology have become more widely adopted in the clinical setting. Four pregnant patients underwent CT scanning with ATCM and were thus further investigated in a pilot study. This includes patient MAM-5 scanned on Philips Brilliance 40 at the first trimester, patient MAM-16 scanned on Siemens Definition AS at the first trimester, and patients MAM-18 and MAM-28 scanned on Siemens SOMATOM Force at the second trimester. An in-house MATLAB code was used to derive the tube current modulation at each slice from the DICOM images. The extracted slice-specific tube currents were used for the calculation of radiation dose at each layer. The fetal organ dose was obtained by summing the radiation doses for each simulated layer.

3 | RESULTS

3.1 | Patient-specific fetal phantom

Figure 3 illustrates the 3D digital constructed maternal phantoms and Table 1 presents the measured anatomical parameters for each patient. The GA ranged from

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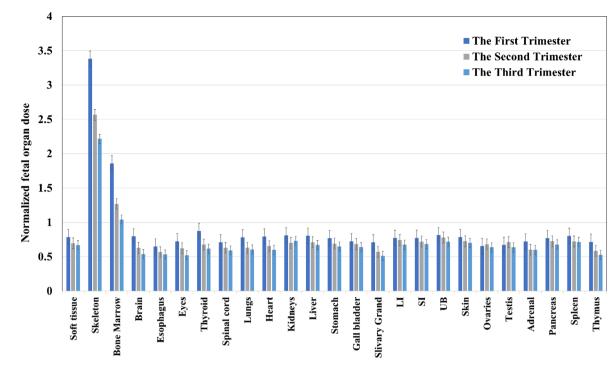


FIGURE 4 Comparison of normalized fetal organ doses (average of three CT scanners and expressed as mean ± SD) at different gestational ages.

8 to 35 weeks. The abdominal circumference of pregnant patients varied between 70.4 and 127.9 cm with a median of 98.27 cm. The scan length of the pregnant woman accepted CT ranges between 23.7 and 64.7 cm. The minimum and maximum FAC were 148.9 and 349.2 mm, respectively. The FL, HL, BPD, and HC were within ranges 27.88-72.68 , 26.01-64.31 , 8.69-97.02, and 32.71-341.54 mm, respectively. The ranges of FD_{skin} were from 19.80 to 84.73 mm, while FD_{skull} varied from 63.22 to 173.90 mm.

3.2 | Fetal organ doses

The fetal organ doses from the three simulated CT scanner systems were estimated and normalized by dividing absorbed dose by the measured CTDI_{vol} of each scanner. Figure 4 shows the calculated normalized fetal organ doses at different trimesters of pregnancy. Consistent with result reported in the literature, it was observed that the fetal organ dose decreases gradually with increasing gestational age.¹⁵ The absorbed doses to the fetal skeleton and bone marrow were about 3.77 and 1.92 times higher than that of fetal soft-tissue. Table 2 shows the correlations between normalized fetal organ dose and the measured anatomical parameters, in which stepwise multiple linear regression analysis was used. As shown in Figure 5, significant correlations were observed between fetal organ dose and MAC, gestational age (GA), and FDs, while we failed to show

statistical correlations between fetal organ dose and FAC, FL, HL, BPD, HC, and scan length. Linear fitting of fetal organ dose was performed for MAC, GA, and FD_{skin} with each point representing one organ in the fetus of an individual patient.

To depict the variability of radiation doses among the three simulated CT scanners, the radiation dose to the fetal brain is compared in Figure 6. It can be observed that the absorbed dose decreases with MAC and the mean absolute dose difference between the developed CT models is 2.78%, thus indicating that the CTDIvol normalized absorbed dose may potentially be less affected by the different CT models than that reported by Turner et al.²⁶ The calculated average fetal organ doses from the three CT systems for skeleton, brain, kidney, liver, and body are shown in Figure 7. For pregnant patients undergoing CT examination, the radiation dose to the fetal skeleton and bone marrow are higher than other organs, whereas the dose to the fetal brain was the lowest. Radiation doses to other organs were found to be similar to each other.

3.3 | Fetal organ dose estimation model

As shown in Table 2, the fetal organ dose demonstrated statistically significant correlations with MAC, GA, and FD_{skin}. The measurements of MAC and GA for individual patients can be routinely obtained in clinical setting

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Fetus Thymus –.871* –.850* –.884* –.612* –.570* –.633* –.862* –.839			839*874*	502*	499*	523*

MEDICAL PHYSICS-

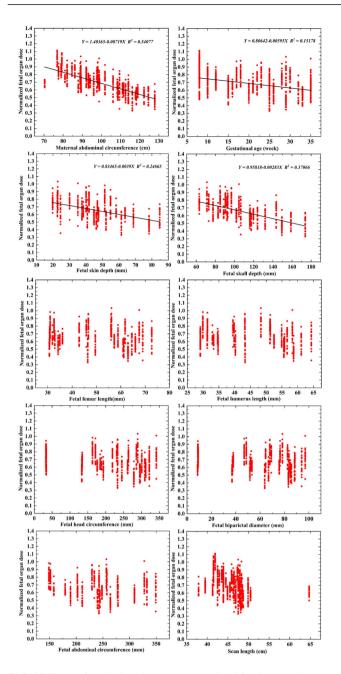
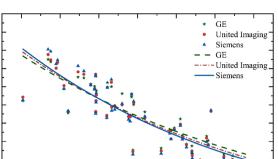


FIGURE 5 Correlations between normalized fetal organ doses and measured anatomical parameters. Stepwise multiple linear regression was performed to analyze the correlation of fetal organ dose with MAC, GA, FD_{skin}, and FD_{skull}. The normalized fetal organ dose was averaged among three CT models.

without access to medical images, whereas FD_{skin} can be measured from ultrasound examinations. From the initial 44 digital patient models investigated, two models incorporating the attainable attributes of MAC, GA, and FD_{skin}, for fetal organ dose estimation in abdominal and pelvic CT imaging were determined:

Model I: Normalized organ dose

$$= \alpha + \beta_1 \times MAC + \beta_2 \times GA \tag{1}$$



1.2

1.1

1.0

0.9

0.8

0.7 0.6

MEDICAL PHYSIC

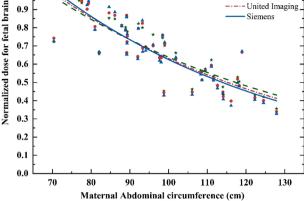


FIGURE 6 Normalized radiation doses to the fetal brain for the three considered CT scanners.

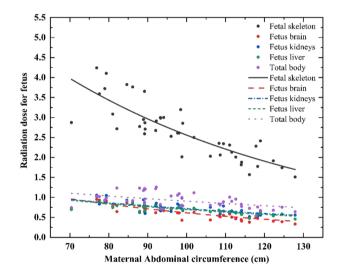


FIGURE 7 The correlations between scanner-averaged normalized fetal organ doses with maternal abdominal circumference of pregnant patients for fetal skeleton, fetal brain, fetal kidney, fetal liver, and fetal body.

Model II: Normalized organ dose

$$= \alpha + \beta_1 \times MAC + \beta_2 \times GA + \beta_3 \times FD_{skin}$$
(2)

where the normalized organ dose refers to the fetal absorbed dose per CTDI_{vol} in abdominal and pelvic CT examinations. Table 3 lists the determined mathematical form of dose estimation models for each fetal organ, while models presenting with $R^2 < 0.5$ were discarded. In most cases, three-factor models yielded better fit than two-factor models as indicated by a higher value of R^2 . For example, the dose estimation formulas of Model II

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for fetal total body can be calculated as:

Normalized organ dose_{II} (fetal total body) = 2.519

$$-0.012 \times MAC - 0.01 \times GA - 0.001 \times FD_{skin}$$
, $R^2 = 0.919$

In specific abdominal CT examinations for pregnant female, the absolute radiation dose (in mGy) to the fetus can be calculated from the calculated normalized organ dose by:

Absolute radiation dose

$$= CTDI_{vol} \times Normalized organ dose$$
 (3)

Where CTDI_{vol} refers to CT scanner's CTDI_{vol} in a specific protocol and can be planned prior to the examination or extracted from the DICOM images in clinical settings.

3.4 | Automatic tube current modulation (ATCM) pilot study—Initial assessment on four cases

The tube current values of each slice are depicted on the red line in Figure 8 where the blue line refers to the average tube current. The normalized fetal dose for these patients from ATCM and FTC examinations were calculated using the developed methodology. That is, the tube current value of each layer was obtained by interpolation, corresponding to simulation calculation settings (just longitudinal (z-) ATCM was accounted for, but no angular tube current modulation), and then the fetal organ dose was calculated. In order to compare the effect of ATCM on fetal dose, the relative deviation between ATCM and FTC was calculated, and the results are shown in Table 4. The tube current is higher than the average value in the scan range of the fetus for patients MAM-16 and MAM-18 as well as the fetal head region of patient MAM-28, which resulted in a higher radiation dose when using ATCM to the fetus of patients MAM-16 and MAM-18 as well as the fetal brain and eyes of patient MAM-28 compared to the FTC protocol. Moreover, the larger size and higher ratio of the visceral belly fat in patients MAM-16 and MAM-18 may result in an increase of tube current when using ATCM at the location of the fetus, leading to increased fetal organ doses.

This study included the largest patients' population covering the whole gestational period using state-ofthe-art CT scanners. The mean absolute fetal dose was 0.72 mGy for the Siemens SOMATOM Force using a low-dose ATCM protocol, 13.87 mGy for the Philips Brilliance 40 using a standard ATCM protocol, 13.73 mGy for the Siemens Definition AS using a standard ATCM protocol, and 1.64 mGy for the GE LightSpeed VCT and **TABLE 3** Normalized organ-level radiation dose estimation models to the fetus with Model I: Normalized organ dose = $\alpha + \beta_1 \times$ MAC + $\beta_2 \times$ GA and Model II: Normalized organ dose = $\alpha + \beta_1 \times$ MAC + $\beta_2 \times$ GA + $\beta_3 \times$ FD_{skin}. (R^2 refers to the coefficient of determination R-squared).

E . (.)		Coeffi	cient			
Fetal tissue/Organs		α	β1	β2	β3	R ²
Brain	Model I	1.536	-0.008	-0.005	_	0.807
	Model II	1.521	-0.007	-0.007	-0.002	0.775
Skeleton	Model I	6.163	-0.299	-0.029	_	0.822
	Model II	5.467	-0.021	-0.025	-0.008	0.894
Bone	Model I	3.354	-0.016	-0.024	_	0.826
marrow	Model II	2.702	-0.010	-0.016	-0.005	0.837
Soft	Model I	1.392	-0.007	0.001	_	0.723
tissue	Model II	1.305	-0.005	-0.001	-0.003	0.849
Esophagus	Model II	1.224	-0.005	-0.002	-0.002	0.813
Eyes	Model I	1.339	-0.007	-0.003	_	0.731
	Model II	1.359	-0.009	0.001	0.001	0.669
Thyroid	Model II	1.566	-0.009	0.000	-0.001	0.802
Spinal cord	Model II	1.360	-0.005	-0.005	-0.003	0.777
Lungs	Model I	1.420	-0.007	-0.002	_	0.737
	Model II	1.266	-0.005	-0.001	-0.003	0.822
Heart	Model I	1.417	-0.007	-0.003	_	0.773
	Model II	1.312	-0.006	-0.002	-0.002	0.785
Kidneys	Model I	1.445	-0.008	0.003	_	0.526
	Model II	1.286	-0.005	0.004	-0.003	0.599
Liver	Model I	1.394	-0.007	0.000	_	0.697
	Model II	1.293	-0.005	0.001	-0.001	0.732
Stomach	Model 1	1.366	-0.007	0.000	_	0.544
Gall	Model I	1.230	-0.006	0.001	_	0.606
bladder	Model II	1.193	-0.005	-0.001	-0.001	0.556
Sliver	Model I	1.348	-0.007	-0.003	_	0.801
Grand	Model II	1.319	-0.007	-0.002	-0.001	0.798
LI	Model 1	1.309	-0.006	0.001	_	0.549
SI	Model I	1.344	-0.007	0.001	-	0.549
	Model II	1.318	-0.005	-0.002	-0.002	0.522
Skin	Model I	1.407	-0.007	0.002	_	0.732
	Model II	1.354	-0.005	0.000	-0.003	0.888
Adrenal	Model I	1.306	-0.007	0.000	_	0.626
	Model II	1.160	-0.004	0.000	-0.003	0.708
Pancreas	Model 1	1.327	-0.007	0.001	-	0.530
Thymus	Model I	1.312	-0.007	-0.003	_	0.801
	Model II	1.249	-0.006	-0.002	-0.001	0.813
Other organs ^a	Model I	1.371	-0.007	-0.001	-	0.543
Total	Model I	1.716	-0.010	0.012	-	0.532
body	Model II	2.519	-0.012	-0.010	-0.001	0.919

^aOther organs are all organs of the fetus except the skeleton and bone marrow.

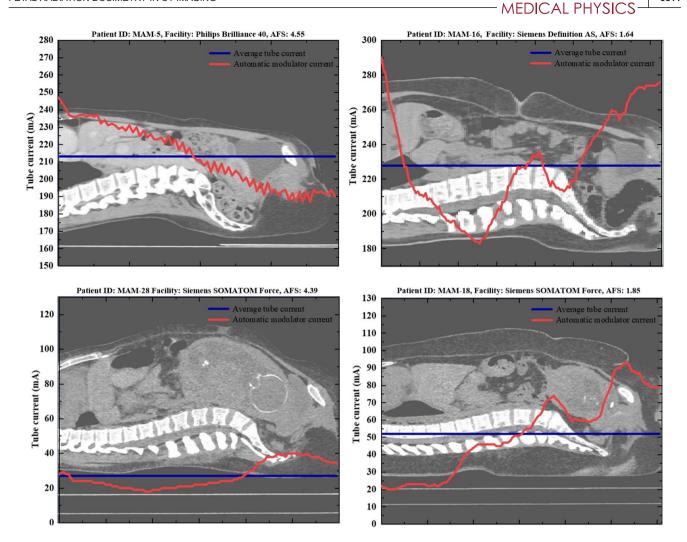


FIGURE 8 Automatic tube current modulation in real-world abdominal and pelvic CT examinations for the studied pregnant patients.

GE Discovery CT750 HD using a low-dose FTC protocol, respectively.

4 DISCUSSION

In this work, three CT scanner systems were utilized for the estimation of fetal dose in 44 pregnant patients undergoing abdominal/pelvic CT examinations. The average fetal radiation dose per 100 mAs from the FTC protocol was determined to be 7.16 mGy/100 mAs within the range of 4.24-9.81 mGy/100 mAs. The absolute fetal radiation dose varied from 0.72 to 13.87 mGy across different imaging protocols for the investigated pregnant patients at ATCM. Since the fetus has high radiosensitivity and conceptus radiation dose is correlated with childhood cancer after birth, accurate and clinically feasible estimation of fetal organ absorbed doses are crucial in radiological imaging of pregnant female patients. The existing studies mostly reported the radiation dose to fetal total body based on simplified fetus model. In this work, we (i) constructed individual fetus models for each pregnant patient based on corresponding medical images, (ii) calculated fetal organ radiation dose using three CT scanner systems, (iii) analyzed the relationship between the fetal organ dose and various anatomical parameters, and (iv) proposed two computational models for dose estimation of fetal organs from abdominal and pelvic CT examinations.

In addition, we compared the fetal organ dose between various scanning protocols on different CT scanners from the major vendors. We observed that the ATCM technique does not necessarily lead to dose reduction to fetal organs, which we suspect is correlated with the location of fetal organs.

With the use of our newly developed computational models, we found that overall, fetal organ radiation dose decreases with increasing gestational age. As known, in the first trimester, a fetus is highly radiosensitive and would have the highest absorbed dose from CT examinations. Our results showed that the normalized fetal organ dose negatively correlated with MAC, GA, and

TABLE 4	Comparison of organ dose between automatic tube	Э
current modu	ation and fixed tube current for pregnant patients.	

Tissue/	Patient II)		
Organs	MAM-5	MAM-16	MAM-18	MAM-28
Fetal soft tissue	-4.44%	7.84%	24.49%	-2.63%
Fetal skeleton	-4.02%	8.15%	26.26%	11.52%
Fetus Bone Marrow	-4.59%	7.66%	27.40%	18.67%
Fetus Brain	-4.76%	7.95%	29.46%	26.54%
Fetus Eyes	-4.08%	8.65%	28.78%	18.44%
Fetus Lungs	-4.34%	7.33%	23.51%	2.29%
Fetus Heart	-4.19%	7.91%	24.30%	2.83%
Fetus Kidneys	-3.98%	7.29%	22.06%	-9.04%
Fetus Liver	-3.94%	7.54%	22.50%	-7.15%
Fetus Stomach	-4.10%	8.20%	23.71%	-7.75%
Fetus Gall bladder	-3.95%	7.44%	21.61%	-11.32%
Fetus Salivary Grand	-4.31%	8.30%	25.59%	16.48%
Fetus LI	-3.77%	8.12%	23.08%	-12.07%
Fetus SI	-3.89%	8.11%	22.79%	-11.72%
Fetus UB	-3.63%	7.80%	22.57%	-13.90%
Fetus Adrenal	-4.05%	7.47%	22.26%	-6.85%
Fetus Pancreas	-3.80%	7.61%	22.54%	-12.29%
Fetus Spleen	3.15%	8.66%	23.91%	-8.78%
Fetus Thymus	17.81%	8.07%	24.36%	6.40%
Average	-2.57%	7.90%	24.27%	-0.02%

FDs, but weak correlation was found with FL, HL, HC, BPD, FAC, and scan length. The absorbed dose to the fetal skeleton was shown to be significantly higher than bone marrow and other organs. This is mainly due to the higher density of the skeleton, which results in the absorption of more scattered radiation. Therefore, prior to performing radiological imaging procedures on pregnant females, the radiologist may give special consideration to the absorbed dose and radiation risks to the fetal skeleton and bone marrow, especially in the first trimester.

The effect of ATCM on fetal radiation dose is rarely discussed and is often overlooked. In our study, the tube current modulation schemes of ATCM extracted from DICOM images of various pregnant patients from different CT scanners were reported. In some cases, ATCM did not improve dose savings to the fetus in pregnant patients and even worse, leads to increased fetal organ dose. For patients MAM-16 and MAM-18, the radiation dose to the fetus when using the ATCM protocol was higher than the one without the FTC protocol. ATCM is aimed to adjust the tube current according to the attenuation characteristics of the patient on the premise of ensuring good image quality as well as reducing the radiation exposure.²⁷ It has been reported that with the increase of cross-sectional dimensions of patients, the tube current in ATCM will increase accordingly to

ensure the quality of CT images and may result in the increase of absorbed dose.²⁸ Hence, we compared the abdominal size of these patients by measuring the abdominal fat ratio (AFS) values,²⁹ which refers to the distance from the anterior spine to the anterior abdominal wall divided by the maximum distance from the outer edge of the rectus sheath to the dermis in the same CT image. To some extent, AFS also indicates the fat layer thickness/body circumference of a pregnant woman during pregnancy. The calculated AFS values for patients MAM-5, MAM-16, MAM-18, and MAM-28 were 4.55, 1.64, 1.85, and 4.39, respectively. The lower AFS for MAM-16 and MAM-18 indicated the higher rate of visceral belly fat and may result in an increase of tube current and absorbed radiation dose to the fetus when using the ATCM protocol.

Table 5 compares the reported fetal radiation dose from CT examinations between this study and the existing literatures^{4,6,10,15,21-25} along with the current modulation mode (ATCM/FTC), patient number, CT protocol number, and gestational age. As shown in Table 5, the calculated fetal doses from this work were lower than the values reported by Hardy et al.,¹⁰ which can be mainly attributed to the differences in CT scanners and the computational pregnant female phantoms. The absorbed fetal dose to patients undergoing CT examinations using the Siemens SOMATOM Force was found to be as low as 0.72 ± 0.81 mGy, which is due to the low tube current and CTDI_{vol} of the chosen protocol. For FTC in two GE CT scanners, the absolute fetal radiation dose was found to be lower than that reported in the literature, which can be attributed to the use of a low-dose scanning protocol, different CT scanners, and individual differences among pregnant patient models. The fetal absorbed dose for all investigated patients in this study were shown to be far below the minimum negligible risk level of 50 mGy for pregnant female receiving a single radiological diagnostic imaging procedure. However, when repeated standard dose CT examinations are needed for pregnant patients in the clinical setting, the accumulated fetal dose may lead to non-negligible radiation hazard and cancer risks for the developing fetus and after-birth child.

This study inherently bears some limitations. Due to the use of low-dose scanning protocols for pregnant patients, most of the fetal internal soft-tissues are difficult to segment within the CT images. The approximation of using the developed anchor phantom and the reference fetal developing measurements provided by the WHO and ICRP may introduce individual uncertainties for radiation dose calculations to these organs. This is a pilot study focusing on an initial assessment of four cases. The patient population evaluated in this work with respect to ATCM needs to be expanded further to include more generalized patterns of CT scanners and scanning protocols. Future work will

TABLE 5 Compo	arison of the	Comparison of the total-body fetal doses between this work and those reported in the literature.	nuses between un		-					
	jo V	No. of real	Range of	Current	Fetal radiation dose (mGy)	e (mGy)	Fetal radiation dose per 100 mAs (mGy/100 mAs)	dose per 100 mAs)	Fetal radiation dose per CTDI _{vol}	dose per
	patients		age	mode	Average \pm SD	Range	Average	Range	Average \pm SD	Range
This study	44	5/3	8~35 weeks	ATCM ^f	$0.72 \pm 0.81^{a},$ $13.87^{b}, 13.73^{c}$	0.39~13.87	7.38 ± 2.61	6.54~9.11	0.95 ± 0.49	0.84~1.17
				FTC	$1.64 \pm 0.33^{d,e}$	$0.97 \sim 2.24$	7.16 ± 1.66	4.24~9.81	0.95 ± 0.25	$0.56 \sim 1.30$
Hardy et al. ^{10,21}	24	6/1	$5\sim36$ weeks	ATCM ⁹	16.61 ± 4.10	10.86~22.39	I	I	1.43 ± 0.24	1.09~1.87
				FTC	21.30 ± 5.26	14.87~28.12	I	I	1.37 ± 0.26	0.96~1.81
Matsunaga et al. ²²	.	0	30 weeks	FTC	2.93 ± 0.41	2.5~3.3	I	I	I	Ι
Xie et al. ⁴	30	1/1	$8{\sim}35$ weeks	FTC	I	Ι	9.20 ± 1.60	6.70~13.0	1.17 ± 0.20	0.85~1.63
Gu et al. ⁶	c	0	3, 6, 9 months	FTC	12.22 ± 8.00	7.81~20.92	6.90	I	I	Ι
Matsunaga et al. ²³	.	2/1	30 weeks	FTC	3.1	I	Ι	I	I	I
Damilakis et al. ²⁴	117	1/1	$0{\sim}7$ weeks	FTC	I	I	Ι	I	0.62 ± 0.29	
Angel et al. ¹⁵	24	6/1	$5\sim36$ weeks	FTC	I	I	11.06 ± 2.24	9.51~14.52	I	I
Felmlee et al. ²⁵	. 	4/3	I	FTC	I	I	11.30	9.00~13.60	I	I
^a Siemens SOMATOM Force: collimator = 57.6 mm, tube voltage = 120 kVp, mean effectiv ^b Philips Brilliance 40: collimator = 40 mm, tube voltage = 120 kVp, mean effective mAs = ^c Siemens Definition AS: collimator = 19.2 mm, tube voltage = 120 kVp, mean effective mAs = ^d GE LightSpeed VCT: collimator = 40 mm, tube voltage = 120 kVp, mean effective mAs = ^e GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective m ^g GE Discovery CT750 HD: collimato	orce: collimat olimator = 40 ollimator = 40 HD: collimator = 40 HD: collimator irrent modifica	ior = 57.6 mm, tube mm, tube voltage = 19.2 mm, tube volta 19.2 mm, tube voltage : = 40 mm, tube voltage : = ation extracted from patients, "0" represe	voltage = 120 kVp, r = 120 kVp, mean effe gge = 120 kVp, mean efft age = 120 kVp, mean efft age = 120 kVp, mean age = 120 kVp, mean i DICOM header of p	mean effective mAs scrive mAs = 162.51 a effective mAs = 12 ective mAs = 22.9,t in effective mAs = 2 patient.	^a Stemens SOMATOM Force: collimator = 57.6 mm, tube voltage = 120 kVp, mean effective mAs = 22.17 (MAM-28) / 43.03 (MAM-18), table feed per rotation = 34.5 mm/s, pitch = 0.60 ^b Philips Brilliance 40: collimator = 40 mm, tube voltage = 120 kVp, mean effective mAs = 162.51, table feed per rotation = 52.5 mm/s, pitch = 0.656. ^c Stemens Definition AS: collimator = 19.2 mm, tube voltage = 120 kVp, mean effective mAs = 141.93, table feed per rotation = 52.5 mm/s, pitch = 0.656. ^d GE LightSpeed VCT: collimator = 40 mm, tube voltage = 120 kVp, mean effective mAs = 141.93, table feed per rotation = 15.4 mm/s, pitch = 0.80. ^d GE LightSpeed VCT: collimator = 40 mm, tube voltage = 120 kVp, mean effective mAs = 22.9, table feed per rotation = 55 mm/s, pitch = 1.375. ^d GE Discovery CT750 HD: collimator = 40 mm, tube voltage = 120 kVp, mean effective mAs = 22.8, table feed per rotation = 55 mm/s, pitch = 1.375. ^g Patient-specific tube current modification extracted from DICOM header of patient. ^m As = 22.88, table feed per rotation = 55 mm/s, pitch = 1.375. ^f Patient-specific tube current modification extracted from DICOM header of patient.	(MAM-18), table fer 52.5 mm/s, pitch = (ion = 15.4 mm/s, pitc 55 mm/s, pitch = 1.37 on = 55 mm/s, pitch =	ad per rotation = 34. .656. h = 0.80. 5. = 1.375.	5 mm/s, pitch = 0.6	0.	

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report on the generation of tube current modification pattern of the ATCM protocol to reduce fetal radiation dose.

5 | CONCLUSION

A clinically feasible framework for calculation of fetal organ-level radiation dose from abdominal and pelvic CT examinations was developed based on personalized computational phantoms of a series of fetus representative of a comprehensive range of gestational age, maternal size, and fetal anatomical characteristics. Fetal organ radiation doses were calculated for three CT scanner systems using validated Monte Carlo simulations. The correlation between fetal organ doses and anatomical parameters, fetal developing indicators, and clinical scanning protocols was investigated. The use of ATCM for abdominal and pelvic CT scanning on pregnant patients was also investigated indicating that it may not lead to radiation dose savings for the fetus.

ACKNOWLEDGMENTS

This work was supported by National Natural Science Foundation of China (32227801) and the Swiss National Science Foundation under grant SNSF 320030_176052.

CONFLICT OF INTEREST STATEMENT

The authors declare no conflicts of interest.

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SUPPORTING INFORMATION

Additional supporting information can be found online in the Supporting Information section at the end of this article. **How to cite this article:** Qu S, Liu H, Xie T, Giger ML, Quan G, Zaidi H. Patient-specific fetal radiation dosimetry for pregnant patients undergoing abdominal and pelvic CT imaging. *Med Phys.* 2023;50:3801–3815. https://doi.org/10.1002/mp.16304