Influence of risk-organ–based tube current modulation on CT-induced DNA double-strand breaks in a biological phantom model

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ABSTRACT
Techniques for dose reduction in computed tomography (CT) are receiving increasing attention. Lowering the tube current in front of the patient, known as risk-organ–based tube current modulation (RTM), represents a new approach. Physical dose parameters can determine the exposure but are not able to assess the biological–X-ray interactions. The purpose of this study was to establish a biological phantom model to evaluate the effect of RTM on X-ray–induced DNA double-strand breaks (DSBs). In breast phantoms and in the location of the spine in an Alderson phantom, isolated human blood lymphocytes were irradiated using a 128-slice CT scanner. A standard thoracic CT protocol (120 kV, 110 ref. mAs, anatomy-based tube current modulation, pitch 0.6, scan length 30 cm) with and without RTM was used. X-ray–induced DSBs were quantified in isolated blood lymphocytes using immunoﬂuorescence microscopy after staining for the phosphorylated histone variant γ-H2AX.

Using RTM, the resulting DNA damage reduction was 41% in superficial breast locations (P = 0.0001), 28% in middle breast locations (P = 0.0003) and 29% in lower breast locations (P = 0.0001), but we found a DNA damage increase of 36% in superficial spine locations (P = 0.0001) and of 26% in deep spine locations (P = 0.0001). In summary, we established a biological phantom model that is suitable for detecting DNA damage in distinct organs. In addition, we were able to show that, using RTM, X-ray–induced DNA damage in the breast can be significantly reduced; however, there is a significant increase in DSBs in the location of the spine.

Keywords: DNA double-strand breaks (DSBs); γ-H2AX; risk-organ–based tube current modulation (RTM); computed tomography (CT)

INTRODUCTION
Computed tomography (CT) contributes the major proportion to radiation exposure in diagnostic imaging [1–3]. Therefore, techniques for dose reduction are receiving increasing attention. Several techniques (like anatomy-based tube current modulation, low-kV protocols, and—particularly used in cardiac CT—ECG-triggered tube current modulation and high-pitch protocols) are already in use [4–6]. Risk-organ–based tube current modulation (RTM) is a rather new approach in CT, and it is supposed to reduce the dose to radiation-sensitive risk organs such as the female breast. RTM is based on lowering the tube current while the X-ray tube is in front of the patient, but increasing it when it is to the rear of the patient to maintain image quality. Estimation of the delivered radiation dose during CT is based on physical measurements using ionization chambers, mathematical analyses (e.g. Monte-Carlo simulation) and calculation of exposure parameters like computed tomography dose index (CTDI) or dose length product (DLP) [1, 7–9]. However, the biological radiation effects not only depend on the radiation dose, but are also reliant on individual patient factors like age and DNA repair capacity, and the application of contrast medium [4, 6, 9–15]. γ-H2AX immunofluorescence microscopy is a sensitive technique for determining X-ray–induced DNA double-strand breaks.
breaks (DSB) in vitro and in vivo. Previous studies demonstrated a very good correlation between radiation dose and the resulting γ-H2AX foci representing distinct DSBs in human blood lymphocytes after CT, angiography or cardiac catheter examination [12, 16–19]. However, in those studies peripheral blood lymphocytes were used to assess radiation damages; that technique, however, is not suitable for evaluation of the effect of RTM because in this case the radiation effects on the organ are more of interest than the effects on the whole body. Therefore, the aim of this study was to establish a biological phantom model for estimation of the X-ray–induced γ-H2AX foci in radiation-sensitive organs, and to evaluate the effect of RTM.

MATERIALS AND METHODS
The study complies with the Declaration of Helsinki and was performed following local ethics committee approval. Written informed consent was obtained from every volunteer.

Preparation of blood samples
For the in vitro experiments, blood samples from three healthy volunteers (median age: 33.2 years, range 27–38 years) with no history of cancer, chemotherapy or leukemia and without exposure to ionizing radiation within the last 3 days were used. Thirty milliliters of blood was obtained from the antecubital vein of each individual using ethylenediaminetetraacetic acid (EDTA)-containing vials.

To isolate human blood lymphocytes, blood was layered onto lymphocyte separation medium 1077 (Biochrom, Berlin, Germany) and centrifuged at 1200g for 15 min at a temperature of 37°C. Cells from the resulting interphase containing ~80% lymphocytes [17] were resuspended in a solution containing 80% Roswell Park Memorial Institute (RPMI) medium 1640 (Biochrom, Berlin, Germany) and 20% fetal calf serum (FCS). These cells were washed and centrifuged twice at 300g and once at 240g, respectively for 10 min, resuspended in phosphate buffered saline (PBS) (ph 7.1) and distributed into small (2 μl) plastic tubes. The separation and washing processes lasted ~50–60 min and were performed at 37°C.

Phantom model and CT protocols
For the exposure, a 128-slice CT-Scanner (Siemens Definition Flash, Siemens Healthcare, Forchheim, Germany) was used. To establish the phantom model we used a commercially available 32 cm CTDI-phantom. Tubes containing 1 μl of isolated lymphocyte suspension were placed in the central and in the peripheral boreholes of the phantom. We irradiated the lymphocytes using a standard chest CT protocol (tube voltage 120 kV, pitch 0.6, rotation time 0.28 s) at different tube currents of 50, 100, 150, 200 and 250 mAs without anatomy-based or risk-organ–based tube current modulation. The scan range was 10 cm. This experiment was performed in triplicate.

For evaluation of the risk-organ–based tube current modulation (RTM), an Alderson chest phantom was positioned precisely at a defined height (center of the gantry) by using the localization system of the CT scanner with commercially available breast essays. The breast phantoms consist of three slices of 2.5 cm thickness, each with several boreholes (Fig. 1a and b). Tubes containing 1 μl of lymphocyte suspension were placed in various superficial (Locations 1–4), middle (Locations 5–6) and deep locations (Locations 7–10) in the breast phantoms (Fig. 1a). To evaluate the effect of RTM on the dorsal parts of the body, lymphocyte-containing tubes were placed in three consecutive slices in deep (Locations R1, R3 and R5) and superficial spine locations (Locations R2, R4 and R6) at the same height as the breast locations (Fig. 1b). These experiments were also performed in triplicate.

Non-irradiated lymphocytes were used as controls to determine the γ-H2AX baseline levels (n = 3). The lymphocytes were exposed to X-rays using a standard chest CT protocol as shown in Table 1, either with or without RTM (X-Care, Siemens Healthcare, Forchheim, Germany). The scan range was 30 cm. Using RTM, the tube current was reduced ~40% at an angle of 130° when the tube was in front of the patient. To maintain the image quality, the tube current was increased while the tube was to the rear of the patient. In every experiment, the standard anatomy-based tube current modulation (Care Dose 4D, Siemens Healthcare, Forchheim, Germany) was used. CTDIvol and DLP were registered as provided by the patient protocol (Table 2).

Immunofluorescent work-up of the exposed blood lymphocytes
At 5 min after irradiation, samples were stored at 4°C during transport to the laboratory and work-up was performed immediately. Lymphocyte samples were layered onto microscope slides for 10 min at room temperature, followed by fixation in 100% methanol (20 min, at ~20°C) and permeabilization in 100% acetone (1 min, at ~20°C). All microscope slides were washed three times in PBS containing 1% FCS for 10 min. The fixation process lasted ~40 min.

To quantify the DNA DSBs, staining of the early phosphorylated histone variant H2AX was used. After overnight incubation with a specific γ-H2AX antibody (Anti-H2AX-Phosphorylated (Ser 139), BioLegend, Uithoorn, The Netherlands) at a concentration of 1:2500 at 4°C, and after washing (3 × 10 min) with PBS containing 1% FCS, the microscope slides were stored for fixation in 2.5% formaldehyde for 20 min (~20°C). Each sample was washed three times for 10 min in PBS containing 1% FCS and incubated with Alexa Fluor 488-conjugated goat antinouse secondary antibody (Invitrogen, Paisley, UK) at a dilution of 1:400 for 1 h at room temperature. The lymphocytes were washed 4 × 10 min in PBS (pH 7.1) and mounted with VECTASHIELD® mounting medium containing 4,6-diamidino-2-phenylindole (Vector Laboratories, Burlingame, USA).

γ-H2AX immunofluorescence microscopy
Fluorescence analysis was performed using an Axiosplan 2 microscope (Carl Zeiss, Jena, Germany) equipped with a x63 magnification objective. In each sample, lymphocytes were counted until 40 foci were detected, and the γ-H2AX foci numbers were related to the number of enumerated cells (each γ-H2AX focus represents one DSB). To determine the number of X-ray-induced DNA DSBs (excess foci), pre-exposure foci levels were subtracted from the number of foci obtained after irradiation. The foci of each microscope slide were quantified three times, and each microscope slide
was evaluated independently by two blinded persons with 6 and 11 years of experience in γ-H2AX immunofluorescence microscopy.

**Statistical analysis**

Statistical analyses were performed using the software Prism 4.03, 2005 (Graph-Pad Software, San Diego, CA). The Spearman correlation was calculated between excess foci levels and the tube current time product. In order to compare DSB levels in different locations in the Alderson phantom with and without RTM, the paired t-test was used. A P-value <0.05 was considered to be statistically significant.

**RESULTS**

The pre-exposure mean γ-H2AX foci levels in the blood samples were 0.077 DSBs/cell ± 0.015 and were used to determine the number of X-ray–induced DNA DSBs (excess foci): the pre-exposure
DSBs/cell (50 mAs) to 0.176 DSBs/cell (250 mAs) in the central, with central samples. The mean excess foci ranged from 0.037
Phantom, but the increase in foci was higher in peripheral compared
exposure
time product (Fig. 2).

There was only a non-significant, slight increase in the DLP
using RTM (Table 2). Regardless of the locations in the breast of
Alderson Phantom we detected significantly fewer excess foci when using RTM in all samples compared with in samples exposed
without RTM (see Figs 3a and 4a); thereby, we also observed a
dependence of excess foci from the sample location in the breast
phantoms. The means of the excess foci were 0.165 DSBs/cell ±
0.026 without RTM and 0.098 DSBs/cell ± 0.013 with RTM (P =
0.0001) in superficial locations (1–4), 0.106 DSBs/cell ± 0.014/cell
without RTM and 0.076 DSBs/cell ± 0.008 with RTM (P = 0.0003)
in middle locations (5–6), and 0.059 DSBs/cell ± 0.009 without RTM
and 0.042 DSBs/cell ± 0.008 with RTM (P = 0.0001) in deep
locations (7–10) (Fig. 3a and left panel of Fig. 4a). The resulting DNA
damage reduction was 41% in superficial locations, 28% in middle
locations and 29% in lower locations (see left panel of Fig. 4b).
Furthermore, mean excess foci were 0.126 DSBs/cell ± 0.014 without
RTM and 0.172 DSBs/cell ± 0.014 with RTM (P = 0.0001) in
superficial spine locations (R2, R4 and R6) and 0.078 DSBs/cell ± 0.014
without RTM and 0.098 DSBs/cell ± 0.013 with RTM (P = 0.0001)
deep spine locations (R1, R3 and R5) (see Fig. 3b + 4a right panel).
Using RTM, we obtained a DNA damage increase of 36% in superficial
spine locations (R2, R4 and R6) and of 26% in deep spine locations
(R1, R3 and R5) (Fig. 4b right panel).

**DISCUSSION**

We established a biological in vitro phantom model for estimation
of CT-induced DSBs in distinct organs. Independently of the use of
RTM, we detected the highest foci levels in superficial locations,
followed by middle and deep locations, which can be explained by
absorption of X-rays in the breast tissue. With RTM, a DSB reduc-
tion could be obtained in superficial locations (−41%), followed by
middle and deep locations (−30%). Compared with a standard thor-
acic CT, an increase in DSBs was found in superficial (36.5%) and
deep spine locations (26%), due to an increase in the tube current

### Table 1. Scan protocols: this table shows the scan parameters of the standard chest CT protocol without and with RTM

| Thoracic CT protocol without RTM | Thoracic CT protocol with RTM |
|---------------------------------|--------------------------------|
| Tube current time product (effective mAs) | 110 | 110 |
| Tube voltage (kV) | 120 | 120 |
| Pitch | 0.6 | 0.6 |
| Rotation time (s) | 0.28 | 0.28 |
| Scan time (s) | 4.18 | 4.18 |
| Anatomy-based tube current modulation | + | + |
| Scan length (mm) | 300 | 300 |
| Risk-organ–based tube current modulation | − | + |

### Table 2. Exposure protocols: this table shows the exposure parameters for thoracic CT protocols with and without RTM, as used for the Alderson phantom experiments

| Thoracic CT protocol without RTM | Thoracic CT protocol with RTM |
|---------------------------------|--------------------------------|
| Reference mAs | 110 | 110 |
| Resulting mAs | 95 | 110 |
| CTDIvol (mGy) | 6.44 | 7.51 |
| DLP (mGy × cm) | 214 | 227 |

foci levels were subtracted from the number of foci obtained after
irradiation.

A significant increase in γ-H2AX foci was obtained after X-ray
exposure—regardless of central or peripheral location in the CTDI-
Phantom, but the increase in foci was higher in peripheral compared
with central samples. The mean excess foci ranged from 0.037
DSBs/cell (50 mAs) to 0.176 DSBs/cell (250 mAs) in the central,
and from 0.046 DSBs/cell (50 mAs) to 0.220 DSBs/cell (250 mAs)
in the peripheral samples. Both for central (r = 0.9906; P = 0.0011)
and peripheral locations (r = 0.9911, P = 0.0010), a significant cor-
relation was obtained between the excess foci and the tube current
time product (Fig. 2).

- Fig. 2. Set-up for establishment of the phantom model using the
CTDI phantom: a standard chest CT protocol with a
constant tube voltage of 120 kV and various tube current
time products without anatomy-based or risk-organ–based
tube current modulation. The x-axis shows the tube current
time product, the y-axis represent the excess foci levels. The Pearson
correlation (r), both for the peripheral (squares) and the central locations (circles) are shown. The figure illustrates the mean excess foci of three independent
measurements. A P-value < 0.05 was considered statistically
significant.
to the rear of the patient. This increase can be explained by the fact
that there is an increase in the tube current when the tube is behind
the patient. This is necessary in order to maintain image quality.
Overall, higher DSB levels were obtained in superficial than in deep
locations, which can be explained by X-ray absorption.

In various previous studies, a similar decrease in the radiation
dose by RTM was reported but, in contrast, we are the first group
that has used a biological model. Vollmar et al. studied the effect of
organ-based–risk tube current modulation using the Monte Carlo
simulation and reported a dose reduction of 47.8% in the chest and
an increase in the bone marrow of 17.9% [9]. Duan et al. observed
similar effects with an anthropomorphic phantom. Using an ioniza-
tion chamber, they determined a dose reduction of 38.1% for the
anterior region and an increase of 24.7 % for the posterior parts of
the body [20]. Finally Wang et al. found a dose reduction of
34–39% for the risk organ–based tube current modulation using a
semianthropomorphic phantom [21]. Our results are in line with
those studies, but there has been a recent study by Franck et al. that
showed there is only a 9% dose reduction in the female breast, in
contrast to our results [22]. This difference can be explained in
that, in their study, not all breast tissue was within the reduced tube
current zone, and therefore the effect was not as big as if the breast
tissue was within the reduced tube current zone. Interestingly, the
increase in the radiation dose in the spine location was similar to
our increase in DSBs in the spine location (26% vs 26–36.5%); this
can be explained because, compared with the female breast, the
spine has less anatomical variants in size and shape. Fu et al. showed
that depending on the female breast location, there are differing
amounts of radiation exposure: if the breast tissue is within the
reduced tube current zone, then a dose reduction of ~40% was
detected, which is in line with our results [23].

The immunofluorescence microscopy method, based on the
phosphorylation of the histone variant H2AX, can be used for quan-
tification of DNA DSBs during various radiologic techniques,
including CT and angiography, and it is well established [4, 6,
11–14, 17–19]. A significant correlation between the numbers of

Fig. 3. Excess foci in all of the distinct tested sample locations in the Alderson phantom: (a) shows the values of the breast
phantoms, locations correspond to the locations of Fig. 1a; (b) demonstrates values in the spine locations in three
consecutive slices of the chest phantom (R1, R3 and R5: deep position; R2, R4 and R6: superficial position). The black
columns show samples irradiated using the standard protocol without RTM; the grey columns indicate samples irradiated
using RTM. Columns represent mean excess foci; error bars indicate standard deviation.
X-ray–induced γ-H2AX foci and the dose deposited in vitro and in vivo has been found \[4, 5, 13, 24, 25\]. Each H2AX focus represents one DSB. DSBs are considered to be the most significant DNA lesions induced by ionizing radiation. Previously, it has been shown in a mouse model that induction of DSBs is comparable between different cell types \[19, 26\]. After X-ray exposure, induced DSBs can be observed within 5 min \[4, 5, 17, 19, 24\]. This method is able to detect DNA damage after low radiation exposure (<1 mGy). Therefore, it provides a reliable and sensitive technique for quantification of acute DNA damage \[4, 6, 10–12, 17, 18\]. However, in all of these studies γ-H2AX foci were evaluated in peripheral blood lymphocytes. This approach would not be appropriate for the evaluation of the effect of RTM because this dose-reducing technique, rather than having a systemic effect, has a local effect on the exposed organs. Therefore, we decided to adapt the γ-H2AX immunofluorescence method to estimate the local DNA damage in distinct organs using an Alderson phantom. For the establishment of our phantom model, we performed experiments using a CTDI phantom and obtained a significant correlation between the excess foci levels and the DLP in the tested dose range (\(r = 0.9906; P = 0.0011\)), confirming the reliability of our approach.

There are also some limitations to this study that have to be mentioned. We only performed in vitro experiments, simulating the in vivo situation, because due to ethical reasons and the risk for the patients it is not possible to obtain tissue samples from individuals undergoing thoracic CT. The setting of the study is experimental and the data represent only estimations of the real DNA damage in breast tissue, and we used a standard Alderson phantom, so we cannot describe the effect of the constitution (thin vs obese bodies). In addition, the breast essays of the Alderson phantom do not represent the individual differences in size or amount of glandular tissue. The complete breast tissue is not located within the angle of 130° in all women, the angle within which the tube current is lowered; thus, all breast tissue is not protected by the tube current reduction in every case (e.g. this may not be the case in obese patients); whether a bra can improve this situation should be investigated in further studies.

It is important to note that damage in peripheral blood lymphocytes is not the same as cancer induction. Therefore, γ-H2AX foci after ex vivo irradiation of lymphocytes in peripheral blood may not be taken as a surrogate for cancer induction by radiation exposure, even if increased DSB turnover might statistically be associated with

Fig. 4. Dependency of excess foci on the sample locations in the phantom: Part (a) shows mean excess foci of samples placed in the superficial, middle and deep slices of the breast phantoms, and in the superficial and deep spine locations in the chest phantom. Means (columns) and standard deviation (error bars) of each slice are presented. Part (b) shows the percentage of DSB reduction/elevation in RTM in comparison with the standard thoracic protocol. For this illustration, the values of the samples irradiated without RTM, as shown in panel (a), were considered to be 100%. The black columns show samples irradiated using the standard protocol without RTM; the grey columns indicate samples irradiated using RTM.
an increased rate of DNA misrepair. γ-H2AX immunofluorescence microscopy is a time-consuming and expensive method. No reliable automated scoring system is available so far that avoids a routine analysis in a bigger study population. Finally, we only assessed the effect of RTM on the biological radiation effects, and did not evaluate the image quality because this was beyond the scope of our study. However, in a recent paper, the overall image quality was not significantly different between chest CTs with and without RTM [27].

CONCLUSION

In conclusion, the established biological phantom model is suitable for estimation of X-ray–induced γ-H2AX foci representing DNA DSBs in organs. Using this approach, we were able to show a significant DSB reduction in breast locations but also a significant increase in DSBs in spine locations using RTM. With the established phantom model, the effect of other dose reduction tools (e.g. bismuth shields) or scan modes can be studied in future.

CONFLICT OF INTEREST

There are no potential conflicts of interest to disclose.

FUNDING

No funding has been received for the work.

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