

Shading correction for volumetric CT using deep convolutional neural network and adaptive filter

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Background: Shading artifact may lead to CT number inaccuracy, image contrast loss and spatial nonuniformity (SNU), which is considered as one of the fundamental limitations for volumetric CT (VCT) application. To correct the shading artifact, a novel approach is proposed using deep learning and an adaptive filter (AF).

Methods: Firstly, we apply the deep convolutional neural network (DCNN) to train a human tissue segmentation model. The trained model is implemented to segment the tissue. According to the general knowledge that CT number of the same human tissue is approximately the same, a template image without shading artifact can be generated using segmentation and then each tissue is filled with the corresponding CT number of a specific tissue. By subtracting the template image from the uncorrected image, the residual image with image detail and shading artifact are generated. The shading artifact is mainly low-frequency signals while the image details are mainly high-frequency signals. Therefore, we proposed an adaptive filter to separate the shading artifact and image details accurately. Finally, the estimated shading artifacts are deleted from the raw image to generate the corrected image.

Results: On the Catphan©504 study, the error of CT number in the corrected image's region of interest (ROI) is reduced from 109 to 11 HU, and the image contrast is increased by a factor of 1.46 on average. On the patient pelvis study, the error of CT number in selected ROI is reduced from 198 to 10 HU. The SNU calculated from the ROIs decreases from 24% to 9% after correction.

Conclusions: The proposed shading correction method using DCNN and AF may find a useful application in future clinical practice.

Keywords: Shading artifact, deep convolution neural network; adaptive filter (AF); volumetric CT (VCT)

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Introduction

Volumetric CT (VCT) is widely used in image-guided radiation therapy. Due to scatter and beam hardening in VCT scanning, a low-frequency shading artifact severely deteriorates the quality of the reconstructed image. Shading artifact causes reconstructed image inaccuracy and spatial non-uniformity (SNU). Without shading correction, the CT number error of reconstructed image CT will be exceeded 350 HU, which brings errors to the positioning accuracy of image-guided treatment and the diagnosis of the image. Thus, it limits the VCT in the clinical application. Therefore, shading correction in VCT is one of the most important problems to be solved for improving VCT image

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quality.

The importance of shading correction on the VCT system is reported by many publications on this topic (1-12). The existing methods of shading correction are mainly divided into two types: pre-processing and post-processing. The pre-processing method corrects the shading artifact in VCT image mainly using an additional hardware device, aiming to prevent the scattering photon from reaching the detector. Following are two typical methods for shading correction in pre-processing. The first one is increasing the air gap between object and detector (13). As the air gap widens, the detection efficiency of the scattered photons will be reduced because of the small cone angle. However, this method is limited by the physical space of the VCT system because the spatial distance cannot be increased indefinitely.

Meanwhile, the X-ray dose will be increased to compensate for the enlarged distance, which is not practical in the clinic. The second one is using the anti-scatter grid (14), which can block the scattered signal of incident angle, while the attenuation efficiency of scattered light using this method is not high enough. At present, the commercial grid can only provide about three times reduction rate of the scatter-primary ratio (SPR), which cannot guarantee the quality of VCT image in a high scattering environment. It also needs to increase the radiation dose to compensate for the attenuation of the X-ray intensity.

Although pre-processing can directly prevent the scattering photon from reaching the detector, its limitation is obvious, and post-processing is more practical. Postprocessing methods include analytical modeling (15), Monte Carlo simulation (16,17), modulation method (3), measurement-based method (1-3,6,18), and scatter correction based on prior data (19,20). Analytical modeling method has a fast computation speed, but the accuracy of scattering estimation is not high enough especially in the complex object. Monte Carlo simulation is the "golden standard" for scattering estimation, but the method is timeconsuming. Modulation method adds a high-frequency modulator between kV X-ray tube and the scanned object. According to different response characteristics between primary and scatter signal, they could be separated in the frequency domain. However, the modulator must be static in the projection, which requires a high accuracy geometry VCT system. Due to the gantry rotation wobble, its clinical application is limited. The measurement-based method requires inserted blocker (usually using lead) into the X-ray source and scanned object (1-3,6,21-25). In this way, the detector forms the shadow region that only contains the scatter signal, but such a methodology is difficult to operate by changing the hardware setting of the existing system. Shading correction based on prior data can get the corrected image better, but this method needs additional prior patient information. Therefore, it cannot be used as a general solution for shading correction.

To tackle the issues in shading correction, in this paper, a novel approach is proposed incorporating the deep convolutional neural network (DCNN) and adaptive filter (AF) together to estimate the shading artifact accurately, which fully explore the potential of DCNN in segmentation of the VCT image with severe artifact and extract the shading artifact using AF (26). The proposed method does not depend on the prior image data and is completely compatible with the linear accelerator in image-guided radiotherapy (IGRT). It does not need to change other hardware and scan protocols and simultaneously without increasing scanning time and dose.

Methods

Workflow

As is known, CT numbers of the same human tissues are approximately the same (26). According to this feature, a template image without shading artifact can be constructed by image segmentation and be used as the corrected reference. To isolate the shading artifacts from the raw data, we generate the residual image by subtracting template image from raw data. An AF is applied in the residual image to estimate the shading artifact so that we can extract the shading artifact while maintaining structure and detail.

Figure 1 shows the framework of the shading artifact correction using DCNNAF. Due to the image SNU effect by shading artifact, the conventional segmentation algorithm is difficult to distinguish the different human tissues. We propose a DCNN to extract deep feature about the VCT image with shading artifact so that an accurate segmentation can be achieved to distinguish the different tissues. Before we start the framework, the input VCT images I_{train} and the corresponding labels of segmentation I_{label} are applied to train DCNN. To start the framework, we input the reconstructed VCT images I_0 with shading artifacts to the trained DCNN. The template image I_t is generated from the output of the DCNN. The residual image I_r which is generated by the subtracting segmented image from the uncorrected image has shading artifact,



Figure 1 The framework of shading artifact correction using the deep convolutional neural network and an adaptive filter. The legend in the figure I_0 : raw image; R: residual image; I_i : template image; S^* : shading image; I_c^* : corrected image.

structure and image detail. The structure and image detail is the high-frequency signal while the shading artifact is the low-frequency signal. Therefore, an AF is implemented to the residual image to remove the high-frequency signal. The shading artifact at low frequency can be obtained. Finally, the artifact is compensated for the raw image to get the final image. The following sections introduce the key steps in the workflow, including VCT segmentation using DCNN and shading artifact estimation using an AF.

VCT segmentation using DCNN

Dramatic developments in parallel technology enable the processing of big data for DCNN. Semantic pixelwise segmentation is an ongoing topic of research, which is improved by deep learning. Unlike the existing conventional segmentation algorithm, segmentation using DCNN is not just based on the image intensity, so that it is especially suitable for shading contaminated VCT image segmentation. This paper proposes a deep convolutional encoder-decoder architecture for robust and accurate human tissue segmentation (bone, marrow, muscle, fat, and air), which can generate an accurate template image assisting shading correction.

DCNN architecture

The black dotted box in *Figure 1* demonstrates the DCNN. The encoding part and the decoding part are shown in the left and right side (27), respectively. The encoding part follows the convolutional neural network, and the repeated convolution layers are included. The Rectified Linear Unit (ReLU) layer follows the convolution layer. In the down-sampling step, the max pooling operation with the size of 2×2 and the stride of 2 is applied. The number of feature channels is doubled at each down-sampling step. In the decoding part, a concatenation layer from the correspondingly down-sampling step (shown in the gray arrow in *Figure 1*) is copied to the step of up-sampling. The up-sampling convolutional layer with 2×2 filter kernel contains the half number of feature channels. Therefore, the output image can be resized into the same size as the input image. The 3×3 convolutions with ReLU are followed by the max pooling and up-convolution layers. In the final step, a 1×1 convolution layer and a soft-max layer are proposed to map the target of segmented tissue. The proposed DCNN architecture of tissue segmentation has 24 convolutional layers.

Network training

The DCNN architecture aims to obtain the mapping function between the input image and the output of the segmented image. To train the DCNN is to improve the accurate of pixel classification from the VCT image. The final segmentation output from the DCNN can be written as:

$$Y = F(\Theta, X) = soft \max\left(f_n\left(W_n f_{n-1}\left(\cdots f_2\left(W_2 f_1(W_1 X + b_1) + b_2\right)\cdots\right) + b_n\right)\right) [1]$$

Where X is the input of VCT raw data, Y is the image label with the segmented tissue, b_i is the bias in the i^{th} convolution layer, W_i is the i^{th} convolution layer, f_i is a ReLU function in i^{th} convolution layer, Θ represents all the tuning parameters in the training step. *softmax* is the classification layer, which is widely used in image segmentation. The aim of the DCNN architecture is to figure out a set of optimal parameters Θ with the input of the image to minimize the loss function:

$$\arg\min_{\Theta} L(Y, F(\Theta, X))$$
[2]

In this equation, L represents the loss of cross-entropy in segmentation. Since the loss function and the ReLU are differentiable, the back-propagation algorithm can be applied to minimize Eq. [2]. In this study, the DCNN was trained using the Adam algorithm. The learning rate was initially set at 10^{-3} and the factor for dropping the learning rate is 0.1 in the ten epochs passes. The size of the minibatch was 16. DCNN is implemented using Matlab R2018a on a graphics workstation. It has an Intel Core Xeon E5-2697 v3 CPU and 128 GB RAM. Two GPU cards (Nvidia GTX Titan Xp) are used to accelerate the minimization step of the loss function. The tuning parameters Θ are initialized with a random number between -1 and 1.

Training datasets

In this study, the proposed DCNN architecture is applied to the two datasets for testing the practicability of tissue segmentation. The two datasets are introduced as follows:

(I) Catphan©504 phantom dataset. We have collected ten subjects of the projections with different current and voltage of the X-ray tube. The VCT projections were acquired with tabletop VCT systems. The ten subjects of VCT projections were reconstructed using FDK reconstruction algorithm. The reconstructed voxel is 512×512×200. The 1246

voxel size of the image is $0.5 \times 0.5 \times 0.5$ mm³. A total of 2,000 two-dimensional (2D) VCT images were obtained.

(II) Patient pelvis image dataset. The patient's VCT data are collected on the On-Board Imager (OBI) which is equipped on Varian Trilogy. The pelvis dataset consists of 15 subjects with a different patient. The reconstructed voxel is 512×512×56. The voxel size of the image is 0.98×0.98×2.5 mm³. A total of 840 2D VCT images were obtained.

The VCT images in both datasets are aligned to the corresponding label with the different segmentation targets. In the phantom dataset, Teflon, Delrin, Acrylic, Low-Density Polystyrene, water, polymethylpentene, and air are contoured as the target of segmentation. In the patient pelvis image dataset, we segment the image into the air, fat, muscle, marrow and bone. Two networks are trained for the phantom and patient pelvis, respectively. To verify the DCNN network, we divided the VCT image datasets into three categories: training dataset, the validation dataset, and the testing dataset. The training dataset is 80% of the VCT images, while the validation dataset is 20% of the VCT images. The testing dataset is 200 slice of the phantom image and 56 slices of patient pelvis image. The images in these three datasets are different.

Shading artifact estimation using an AF

The segmentation image is obtained from the output of the DCNN. The corresponding tissue area is filled with the standard CT number of corresponding X-ray tube voltage, to generate a template image I_{ij} and this image has no image details. By subtracting the raw images from the template image I_{tr} the residual image R is generated. The residual image contains the shading artifact and the image detail. The shading artifacts are mainly low-frequency signals while the image details are mainly high-frequency signals. Therefore, a low pass filter can be used to separate the shading artifact and image details. Conventional low pass filter can eliminate the image detail, but the boundary of the anatomical structure in the residual image R may be filtered out simultaneously, resulting in the severe loss of image contrast. Consequently, the filter should balance image smoothing and edge preserving. In this paper, a L0 norm smoothing filter is applied to the residual image which can achieve smoothing the image detail while preserving the edge of the structure. The objective function of the filter is as follow:

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$$S_{k} = \arg\min\left\{\left\|R - S\right\|_{2} + a_{k} \cdot \left\|\partial_{x}S\right\| + \left|\partial_{y}S\right|\right\|_{0}\right\}$$
[3]

Here *S* is the estimated shading artifact, *R* is the residual image. $\partial_x S$ and $\partial_y S$ are the gradient of shading artifact in *x* and *y* direction, respectively. a_k is a weight directly controlling the degree of smoothing. $||R-S||_2$ is a constraint of the image structure similarity. A discrete counting metric is applied in the objective function. Since the first term is the pixel-wise difference while the second term is global discontinuity statistically, discrete and the traditional gradient optimization methods are incapable of solving this problem. We apply a special alternating optimization strategy with half-quadratic splitting to solve the objective function (28).

The smoothing image S_k is significantly depended on the smoothing parameter a_k . In this paper, we propose an adaptive framework to choose a suitable a_k automatically, which can achieve the shading correction and image structure protection. Since the shading artifact deteriorates the spatial uniformity of the image, the goal of shading correction is to improve the spatial uniformity in the same tissue. To find a correct smoothing parameter a_k , we figure out the solution based on the assumption that CT numbers of the same human tissues are approximately the same. From the above assumption, we can know that the ideal CT image has high spatial uniformity in the specific tissue. In this paper, we use a sharp peak in the image histogram of the specific tissue to represent the spatial uniformity of the image. Therefore, a suitable value of the smoothing parameter a_k is calculated as an optimization model to minimize the objective function. The function is written as follows:

$$a^* = \arg\min\left\{ \left\| I_0 - I_c(a_k) \right\|_2^2 + \lambda \cdot \max\left(hist\left(I_c(a_k)\right)\right) \right\}$$

s.t. $a > 0$ [4]

Where I_0 is the input of the uncorrected image, $I_c(a_k)$ is the output of the corrected image with the smoothing parameter setting at a_k . *hist* is the image histogram in the specific tissue. $||I_0 - I_c(a_k)||_2^2$ is the image fidelity term, which can protect the structure of the output image. $\max(hist(I_c(a_k)))$ is the term of image spatial uniformity. λ is a penalty factor which is set at -10^{-3} empirically.

Eq. [4] is solved using the mesh adaptive direct search (MADS) algorithm. The convergence analysis for the MADS algorithm can be found in Ref. (29). It can achieve automatic smoothing parameter setting instead of

cockamamie tuning. After getting a suitable parameter of the filter, the final corrected image can be obtained using the following formula:

$$I_{c}^{*} = I_{0} + S^{*}$$
[5]

Where S^* is the estimated shading artifact with smoothing parameter setting at a^* .

Pseudocode

In summary, we present the pseudo-code using the DCNNAF algorithm for the shading correction in Table 1. Line 1 sets the DCNN architecture and the training parameters. Line 2 gives the optimization control parameters, including the stopping criteria and initial setting. Line 3 is the training step to minimize the cross-entropy using Adam algorithm. Line 4 is the predicting step that is using the trained network to segment the input image. Line 5 is the generation step of the residual image. Line 6-23is the main loop of generating the AF. In the step of the AF, Line 7 is the smoothing step on the residual image in order to estimate the shading artifact. Line 8 indicates the generation of the temporary corrected image. Lines 9-11 apply a barrier function $\psi(a_k)$ to change the constraint optimization problem into an unconstraint optimization problem. The barrier objective function $G(a_k)$ is solved using MADS algorithm shown in line 12–23. The searching step in Lines 12-15 is implemented to find a new iteration that decreases the objective function in M_k . When the step of searching fails to find the decreased value, the polling step in Lines 16-20 is performed in the current iteration. When the step of polling also fails to find the decreased value, the parameter of mesh-size is decreased. Otherwise, the parameter of mesh-size is increased. To terminate the iterative process, the stopping criteria Δ_{tol} should be smaller than a given threshold after a certain number of iterations (k_{max}) in Lines 21–23. Lines 24–25 are implemented to compensate the estimated shading artifact into the uncorrected image to get the final image.

Evaluation

The DCNNAD method is evaluated using the Catphan©504 phantom and patient pelvis cases. The phantom projection is acquired using the tabletop VCT system at Shenzhen Institutes of Advanced Technology, Chinese Academy of Sciences. The geometry of the tabletop VCT system

matches with the Varian Trilogy OBI. We also obtained the phantom image using the narrow collimation in front of the kV tube (a width of around 10 mm on the detector). In this fan beam equivalent geometry, scatter signals are inherently suppressed, and the resultant images were referred to as "scatter-free" reference images for comparison. After the Catphan©504 phantom study, the pelvis image of the patient is included to evaluate the practicability and robustness of the proposed DCNNAF algorithm. The pelvis data sets are acquired from patients on Varian Trilogy OBI at the Department of Radiation Oncology. For comparison, the corresponding planning CT of a patient is also acquired as the reference image.

Table 2 lists the scanning and reconstruction parameters. We apply the image contrast and SNU as quality metrics in the regions of interest (ROIs) of the image. Scatter artifacts are more prominent around objects with high contrasts. On Catphan©504 phantom study, the image contrast was calculated as:

$$contrast = |\mu_r - \mu_b|$$
[6]

Where μ_r is the mean CT number of VCT image inside ROI and μ_b is the mean CT number of VCT image in the surrounding area. Since the scatter signals cause nonuniformity in the VCT image, the SNU (30) is measured as:

$$SNU = \left| \frac{\overline{HU}_{\text{max}} - \overline{HU}_{\text{min}}}{1000} \right| \times 100\%$$
[7]

Where \overline{HU}_{max} and \overline{HU}_{min} are the maximum and the minimum of the mean CT numbers the selected ROIs, respectively. Five ROIs with the same diameter of 10 pixels (5.0 mm) were selected in the VCT image of the Catphan©504 and patient pelvis data.

Results

Catphan©504 studies

Figure 2 shows the effects of the scatter correction using the proposed scheme on the reconstructed VCT images. Due to the scatter signal in the projection, the reconstruction error is significant as shown in Figure 2A and D. Since the shading correction is implemented using the DCNNAF method, the shading artifacts are suppressed as demonstrated in Figure 2B and E. Figure 2C and F shows the referenced fan-beam CT image. The average CT numbers in the

No.	Algorithm 1	Description							
1	$batchSize = 16$; $learningRate = 0.01$; $nClass = 5$; $epochesNum = 4000$; $\Theta = rand(0.1)$; $layerParam = (shown in Figure 1)$	// Initial training parameters							
2	$\lambda = -1000; \ a_0 = 0.1; \ \Delta_0 = 1; \ \Delta_{tot} = 10^{-4}; \ k_{max} = 500; \ \tau_e = 2; \ \tau_c = \frac{1}{2}$	// Initial adaptive filter parameters							
3	$\Theta^* = \arg\min L(I_{label}, F(\Theta, I_{train}))$	// Training the network							
4	$I_t = F(\Theta^*, I_0)$	// segmentation using the network							
5	$I_r = I_t - I_0$	// Generate the residual image							
6	for $k = 1$; $k \le k_{\max}, k + +$	// Main loop							
7	$S_{k} = \arg\min\left\{\left\ R - S\right\ _{2} + a_{k} \cdot \left\ \left\ \partial_{x}S\right\ + \partial_{y}S\right\ _{0}\right\}$	// Smoothing the residual image							
8	$I_{c}(a_{k}) = I_{0} + S_{k},$	// Generate the correcting image							
9	$g(a_k) = \left\ I_0 - I_c(a_k)\right\ _2^2 + \lambda \cdot \max\left(hist(I_c(a_k))\right)$	// Objective function							
10	$\psi(a_k) = \begin{cases} 0 & \text{if } a_k > 0, \\ +\infty & otherwise \end{cases}$	// Barrier function							
11	$G(a_k) = g(a_k) + \psi(a_k)$	// Barrier objective function							
12	$\boldsymbol{M}_{k} = \left\{\boldsymbol{a}_{k} + \boldsymbol{\Delta}_{k}\boldsymbol{D}_{z}: z \in \boldsymbol{N}^{ \boldsymbol{D} }\right\}$	// Searching step: Evaluate $G(\alpha_k)$ in M_k . If							
13	if $\exists a_{k+1} \in M_k$ such that $G(a_{k+1}) < G(a_k)$	$o(a_{k-1}) < o(a_k)$, expana mesn. Otherwise, do the poll step							
14	$a_{k+1} = a_k + \Delta_k D_z; \ \Delta_{k+1} = \tau_e + \Delta_k \ continue$								
15	else								
16	$P_k = \left\{ a_k + \Delta_k d : d \in D_k \right\}$	// Poll step: Evaluate $G(\alpha_k)$ at P_k . If $G(\alpha_{k-1}) < G(\alpha_k)$, expand mesh. Otherwise, contract mesh.							
17	if $\exists a_{k+1} \in M_k$ such that $G(a_{k+1}) < G(a_k)$								
18	$a_{k+1}=a_k+\Delta_k d;\;\;\Delta_{k+1}= au_e\Delta_k;$								
19	else $\Delta_{k+1} = \tau_c \Delta_k;$								
20	endif								
21	end								
22	if $\Delta_k < \Delta_{tol}$; $a^* = a_k$; $S^* = S_k$; break; endif	// Stopping criteria of MADS							
23	endfor								
24	$I_c^* = I_0 + S^*$	// Generation of the final corrected image							
25	Return I [*] _c								

Table 1 Pseudocode of shading correction method using DCNNAF

selected ROIs from *Figure 2A,B,C* are -40, 58 and 69 HU, respectively. Consequently, the average absolute error of CT numbers is 109 and 11 HU in the uncorrected and corrected image, respectively. It demonstrates the significant improvement of the proposed method. We evaluated the

spatial resolution using the modulation transfer function (MTF). A circular object (the red box in *Figure 2*) is selected for MTF calculation. The 50% of MTF magnitude is 4.77 in the corrected image while the 50% of MTF magnitude is 4.56 in the uncorrected image. After correcting the scatter,

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Table 2 magning and reconstruction parameters of the phantom and patient									
Scan protocol	Catphan©504 study	Patient pelvis study							
Imaging parameters									
Scan mode	Full-fan	Half-fan							
X-ray energy	120 kVp	100 kVp							
X-ray tube current	80 mA	80 mA							
Pulse width	13 ms	23 ms							
Source to detector distance	1,500 mm	1,500 mm							
Source to rotation axis distance	1,000 mm	1,000 mm							
Detector size	400-by-400 mm ² 1,024-by-1,024	400-by-300 mm ² 1,024-by-768							
Rotation circular	360 deg circular	360 deg circular							
Number of views	662	667							

0.98×0.98×2.5 mm³

512-by-512-by-56

aconstruction parameters of the phantom and patient Table 2

Reconstruction parameters

Voxel size Volume size



0.5×0.5×0.5 mm³

512-by-512-by-200

Figure 2 Axial and coronal views of Catphan©504 phantom. (A,D) No correction, (B,E) with shading correction, (C,F) reference (fan-beam CT image). The images in the 1st row are in the axial view and the images in the 2nd row are in the coronal view. The selected uniform ROIs (marked with dashed red circle) in (A) indicate the locations where the average CT numbers are calculated using the optimized blocker. The average CT numbers in the selected ROIs from (A,B,C) are -40, 58 and 69 HU, respectively. In Figure 2B, the dashed white line represents the position where 1D profile in Figure 3 is taken. The averaged contrasts and CT number are calculated inside the contrast rods, which are indicated by the digits and white dashed circles in Figure 2A. The display window is [-250, 250] HU.

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the spatial resolution in the corrected image is higher than the uncorrected image. The 1D profiles of the CT number along with the dashed white line passing through the two high-contrast rods in *Figure 2B* is shown in *Figure 3*. The 1D profiles match well with the reference image using the proposed method.

For further evaluation of the scatter correction performance using the proposed scheme, the average CT numbers and contrasts are calculated for the contrast rods in one of the phantom inserts as indicated in *Figure 2A*. The results are summarized in *Table 3* using a fan-beam CT as the reference. The CT number error is reduced from 206 to 13 HU in the selected ROI with the implementation of the proposed method. The image contrast is increased by a



Figure 3 Comparison of the 1D profiles passing through the two contrast rods as shown in *Figure 2B*.

factor of 1.46 on average.

Since the Catphan©504 is of the regular structure with an almost uniform distribution of CT number, a more challenge evaluation will be presented in the heterogeneous pelvis patient study in the next section.

Patient head studies

For more challenging, a patient pelvis image obtained on the clinical VCT is evaluated using the proposed framework. Figure 4 shows the result of the processing image using DCNNAF; the 1-3th row is the axial, coronal and sagittal view image, respectively. The shading artifact severely deteriorates the image as shown in Figure 4A,E,I, leading to spatial non-uniformity in the pelvis image. Figure 4B,F,7 shows the segmentation image. Although the shading artifact poses a big challenge in the segmentation of different tissue, the proposed DCNN method can achieve high accurate segmentation in bone, marrow, muscle, fat, and air mainly due to the deep feature extraction in the training data. Accurate segmentation makes the directly shading correction in image-domain a possible. Figure 4C, G, K shows the corrected image, which is significantly suppressed the shading artifact presented in the raw images. The corrected image using DCNNAF method is comparable to the registered planning CT is shown in Figure 4D,H,L. For quantitative image quality analysis, the error of CT number is reduced from 198 to 10 HU in the soft tissue region enclosed by the solid red circle in Figure 4A. The SNU is calculated in the five selected ROIs which are shown in Figure 4A. The SNU in the uncorrected and corrected image is reduced from 24% to 9%.

Table 3 Comparison of the average reconstruction value and image contrasts measured on the contrast rods of the Catphan@504 phantom

Properties	ROI	1	2	3	4	5	6	7	Average
CT value	Fan-beam CT	-987	337	958	-985	-182	-90	-39	_
	VCT with proposed correction	-969	329	948	-961	-168	-82	-28	-
	Error of CT number with correction	18	8	10	24	14	8	11	13
	VCT without scatter correction	-727	175	601	-712	-173	-117	-76	_
	Error of CT number without correction	260	-162	-357	273	9	-27	-37	206
	CT# improvement	250	158	347	243	10	31	50	_
Contrast	VCT with proposed correction	1,002	237	834	981	272	190	120	_
	VCT without scatter correction	724	155	554	717	181	128	84	_
	Contrast Increase Ratio	1.38	1.53	1.51	1.37	1.50	1.48	1.43	1.46

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Figure 4 Axial, coronal and sagittal views of the reconstructed patient pelvis image. (A,E,I) No correction; (B,F,J) segmentation image using DCNN; (C,G,K) with shading correction; (D,H,L) the reference image using registered planning CT of the same patient. The images in the 1^{st} row are in the axial view, the images in the 2^{nd} row are in the coronal view and the images in the 3^{rd} row are in the sagittal view. The selected uniform ROI enclosed by the solid red circle in (A) indicates the area where the average CT numbers of the VCT images in the A, C and D are calculated. The SNUs are calculated on the selected five ROIs enclosed by the dashed yellow circles in (A). The display window is [-250, 300] HU.

Discussion

In this work, we investigated the DCNN in the segmentation of VCT image with severe shading artifact and proposed an AF to correct the shading artifact. Contrasted with the conventional shading correction method (31), the proposed DCNNAF method not only achieves high-quality image but also has a high computational efficiency due to the image-domain processing. Though we have tackled several issues in shading correction, the proposed scheme can be further improved. First, although the prediction is high computational efficiency, the proposed DCNN architecture spends a long time for data training, which generally takes one day to get a trained model, while the traditional segmentation does not need to train. In the future, we will improve the computational efficiency of data training by reducing the capacity of the network with high accuracy of segmentation. Second, the DCNN architecture needs large numbers of training data, and the corresponding labels of segmentation are needed to be contoured manually. The example of training data and corresponding label are shown in Figure 5A and B. In Figure 5B, it is evident that the label image which is contoured manually has a sharp edge between the different tissues. The error of the contoured

image is unavoidable. Even so, the prediction of the testing image has a highly accurate result of segmentation, and the sharp edge error is eliminated shown in *Figure 5C* and *D*. It demonstrates that the DCNN can tolerate contoured error.

Different parameters will influence the accuracy of segmentation. The filter size of the convolutional layer needs to be optimized. We determine the optimized value by changing the filter size while keeping the other parameter fixed. *Figure 6* shows the influence of the filter size on the patient dataset. As compared with the filter size of 1, 5, and 7, the proposed filter size 3 achieve higher accuracy at the last epoch. Therefore, we choose 3 as the best filter size.

Currently, we apply the DCNNAF method to only focus on patient pelvis data. We will extend the study on all parts of human VCT image. Regarding the patient pelvis, we focus on four tissues, i.e., bone, marrow, muscle, and fat in this paper. More tissue classification needs to be done in the future to improve the robustness of the proposed method around the whole part of the human VCT images, as well as its practicability to clinical application. Unsupervised learning is developing fast (32), and it also can achieve segmentation method. In this study, the proposed workflow still requires a large amount of training data and



Figure 5 Example of pelvis image for training and testing. (A) Axial view image of the pelvis for training; (B) manual contoured image as a label; (C) axial view image of the pelvis for testing; (D) segmentation using the trained network; Display window is [-270, 280] HU.



Figure 6 Change of the accuracy on the validation set of the patient using different filter sizes.

segmentation labels. In the future, our team would do more searches on unsupervised learning that needs less training data on this problem.

Conclusions

In this study, we propose a robust shading correction method using DCNNAF, which improves the VCT image quality. It does not need to change other hardware and scan protocols. The method also does not increase the scanning time, and the deliver X-ray dose. On the Catphan©504 study, the error of the CT number in the corrected image's ROI is reduced from 109 to 11 HU. On the patient pelvis study, the error of the CT number in the selected ROI is reduced from 198 to 10 HU. Besides the high shading correction efficacy, the proposed method possesses several advantages over other existing shading correction approaches, including no dose or extra scan time, no requirement of prior knowledge, easy implementations and high quality of the corrected images.

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Footnote

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