

Received September 18, 2014, accepted October 3, 2014, date of publication October 17, 2014, date of current version November 3, 2014.

Digital Object Identifier 10.1109/ACCESS.2014.2363367

# A Stationary-Sources and Rotating-Detectors Computed Tomography Architecture for Higher Temporal Resolution and Lower Radiation Dose

GUOHUA CAO<sup>1</sup>, BAODONG LIU<sup>2,3,4</sup>, HAO GONG<sup>1</sup>, HENGYONG YU<sup>2</sup>, (Senior Member, IEEE), AND GE WANG<sup>5</sup>, (Fellow, IEEE)

<sup>1</sup>Virginia Tech-Wake Forest University School of Biomedical Engineering and Sciences, Virginia Polytechnic

Institute and State University, Blacksburg, VA 24061, USA

<sup>2</sup>Virginia Tech-Wake Forest University School of Biomedical Engineering and Sciences, Wake Forest University

Health Sciences, Winston-Salem, NC 27157, USA

<sup>3</sup>Key Laboratory of Nuclear Analysis Techniques, Institute of High Energy Physics, Chinese Academy of Sciences, Beijing 100049, China
<sup>4</sup>Beijing Engineering Research Center of Radiographic Techniques and Equipment, Beijing 100049, China

<sup>5</sup>Department of Biomedical Engineering, Rensselaer Polytechnic Institute, Troy, NY 12180, USA

Corresponding author: G. Cao (ghcao@vt.edu)

The work of G. Cao was supported by the Directorate for Engineering, National Science Foundation under CAREER Award 1351936. The work of G. Cao and H. Gong was supported in part by a seed grant through the Institute for Critical Technology and Applied Science, Virginia Polytechnic Institute and State University (Virginia Tech), Blacksburg, VA 24061, USA, and in part by Dr. Cao's New Faculty Startup Package at Virginia Tech. The work of B. Liu, H. Yu, and G. Wang was supported by the National Institutes of Health/National Institute of Biomedical Imaging and Bioengineering under Grant EB011785.

**ABSTRACT** In current computed tomography (CT) architecture, both X-ray tubes and X-ray detectors are rotated mechanically around an object to collect a sufficient number of projections. This architecture has been shown to not be fast enough for patients with high or irregular heart rates. Furthermore, both X-ray beams and detectors of the current architecture are made wide enough, so that the entire object is covered in the lateral direction without data truncation. Although novel acquisition protocols have recently been developed to reduce a radiation exposure, the high radiation dose from CT imaging remains a heightened public concern (especially for cardiac CT). The current CT architecture is a major bottleneck to further increase the temporal resolution and reduce the radiation dose. To overcome these problems, we present an innovative stationary-sources rotating-detectors CT (SSRD-CT) architecture based on the three stationary distributed X-ray sources and three smaller rotating X-ray detectors. Each distributed X-ray source has  $\sim 100$  distinctive X-ray focal spots, and each detector has a narrower width compared with the conventional CT detectors. The SSRD-CT will have a field-of-view of 200 mm in diameter at isocenter, which is large enough to image many internal organs, including hearts. X-rays from the distributed sources are activated electronically to simulate the mechanical spinning of conventional single-beam X-ray sources with a high speed. The activation of individual X-ray beam will be synchronized to the corresponding rotating detector at the opposite end. Three source-detector chains can work in parallel to acquire three projections simultaneously and improve temporal resolution. Lower full-body radiation dose is expected for the proposed SSRD-CT because X-rays are restricted to irradiate a local smaller region. Taken together, the proposed SSRD-CT architecture will enable  $\leq$ 50-ms temporal resolution and reduce radiation dose significantly.

**INDEX TERMS** Computed tomography, temporal resolution, radiation dose, multisource, interior tomography.

# I. INTRODUCTION

Since its Nobel-prize winning invention in 1970s [1], x-ray computed tomography (CT) has been through dramatic developments, and the performance of CT scanners has been improved tremendously. One major improvement worthy of mention is temporal resolution, which has been improved from 5 min at the beginning to sub-second at this moment [2].

The current highest temporal resolution is achieved by a second-generation dual-source CT (DSCT, e.g. Definition Flash CT from Siemens), providing a temporal resolution up to 75ms [3], [4].

The need for a higher CT temporal resolution stems from an obvious fact that organs within human bodies are constantly moving. In particular, because heart is the fastest moving organ within the human body, CT developments have been largely directed at improving temporal resolution for cardiac CT. A cardiac CT scan with higher temporal resolution can 'freeze' heart motion more effectively, thus it provides sharper images and fewer artifacts. Although it is favorably reviewed in many clinical cardiac CT imaging, the state-of-the-art temporal resolution of 75ms works best only when image acquisition is ECG-gated to the relatively quiet diastolic phase [5]–[9]. To image various cardiac phases from patients with irregular heart rates (e.g. arrhythmia), it was estimated that a temporal resolution down to 50ms would be needed [10], [11].

To achieve a higher temporal resolution, researchers have previously investigated two strategies. The first and more obvious strategy is to reach a faster gantry rotation speed. This has been the primary strategy in action during the last few decades. Many folds of improvement in gantry rotation speed has been achieved since 1980s. Currently, the fastest gantry rotation speed is about 300ms. However, increasing the temporal resolution via an even faster gantry rotation speed appears to be beyond today's mechanical limits. Nowadays, the gravitational force on the gantry can reach 30g, about 10 times higher than the acceleration experienced during the space shuttle take-off [2], [12]. An innovative idea for ultrafast CT is the electron beam CT (EBCT), which employs a stationary target ring and a stationary detector ring. Because EBCT sweeps an electron beam across a stationary anode ring electronically (i.e. without mechanical gantry rotation) [13], it can achieve a temporal resolution of about 30ms.

The second strategy is to acquire multiple projections simultaneously through multiple source-detector imaging chains. This strategy is well known to improve the temporal resolution of CT scanners. For example, the 75ms temporal resolution of the DSCT is achieved by arranging two x-ray tubes and two detectors on a single gantry with a gantry rotation time of 270ms. Previous simulations showed that doubling the number of imaging chains on a given gantry is more efficient to increase the temporal resolution than reducing the gantry rotation time by a factor of 2 [14]. However, as demonstrated by the dynamic spatial reconstructor (DSR) in the 1980s and depending on the actual geometry, the number of imaging chains to prevent adjacent x-ray beams from overlapping [15]–[17].

Another important improvement in CT is reduction in radiation dose. The radiation risk (e.g. cancer) from CT imaging is a growing public concern [18]. In clinical practices, it generally requires adhering to the principle of reducing radiation doses as low as reasonably achievable (ALARA). The dose concern is particularly critical for CT perfusion imaging, where a high radiation dose ( $\sim$ 10mSv) can be accumulated from multiple CT acquisitions [19]. The dose issue will become even more prominent if a higher spatial resolution is required. This is because, to double the spatial resolution in all 3 spatial directions without affecting the contrast-to-noise ratio (CNR), the radiation dose has to be increased by a factor of  $2^4$  (i.e. 16-fold increase) [20]. Although a number of dose reduction techniques have been developed [21]–[25], more works are needed to allay public concern on CT radiation dose.

Aiming to improve temporal resolution and lower radiation dose in this paper we propose a new CT architecture by combining the latest compressive-sensing (CS) based interior tomography [26] and distributed x-ray source technology [27], [28]. The interior tomography technique allows accurate reconstruction of a region-of-interest (ROI) from truncated projections. In contrast to the conventional CT architecture, the detectors for interior tomography do not cover the full transaxial extent of the object. The interior tomography technique can help to reduce detector size, suppress scattering artifacts, and lower radiation dose. Furthermore, radiation doses could be further reduced when interior tomography is combined with the compressivesensing framework [26]. A distributed x-ray source can produce x-rays by extracting electron beams from an array of cathodes and sending each electron beam to a distinctive focal spot on the same target. The cathodes are made of either hot dispenser cathode emitters (DCE's) [27] or cold field emitters such as carbon nanotubes (CNT's) [28]. Electron beams (and hence the corresponding x-ray beams) can be switched on and off electronically by applying the corresponding extraction voltages on the cathodes. By programming these extraction voltages, a scanning x-ray beam can be generated to irradiate an object from different angles, enabling tomographic imaging without mechanical motion [29]. The CNT-based distributed x-ray sources are very attractive because the cold CNT cathodes work at room temperatures and allow compact packaging and faster and more flexible control of x-ray beams [30].

In the rest of this paper, we will first present our new architecture, and then demonstrate its feasibility with numerical analyses and simulation using a clinical cardiac CT dataset. The results will be presented and the rationales for the expected performance will be discussed. Finally, we will discuss the potential limitations and challenges for the implementation of the proposed SSRD-CT architecture.

# **II. THE PROPOSED CT ARCHITECTURE**

The proposed CT architecture is illustrated in Fig. 1. It has three stationary distributed x-ray sources and three rotating detectors. Hereafter, we refer this architecture as stationarysource rotating-detector CT (SSRDCT). Three identical source-detector chains are positioned around an object symmetrically. The key geometrical parameters are listed in Table 1. The source-to-isocenter distance (SID) and the source-to-detector distance (SDD) are 540mm and 950mm, respectively, which are close to most of commercially available CT scanners. The field-of-view (FOV) is at 200mm, which is sufficient to cover the transaxial extent of a typical human heart. This design will allow us to reuse, or use with slight modifications, many existing and mature CT technologies in detectors, gantry, slip rings, etc. The core



**FIGURE 1.** Schematics of the proposed CT architecture with three stationary distributed x-ray sources and three rotating detectors.

TABLE 1. Geometrical parameters of the proposed CT architecture.

Number of Source-Detector Chains	3
Source to Isocenter Distance (mm)	540
Source to Detector Distance (mm)	950
Field of View	200

new elements are the interior tomography and the distributed x-ray sources.

Each distributed x-ray tube will have a multitude of x-ray sources that are distributed equiangularly (relative to the isocenter) in a shared vacuum envelope. During a CT scan, the x-ray sources within an x-ray tube will be sequentially turned on and off from one end to the other. As a result, each x-ray source will give a pulsed radiation, and the switching is done through programming the gate voltage on the corresponding cathodes [27], [30]. Furthermore, the switching of x-ray sources will be synchronized with to the rotation of the detectors to simulate the spinning of three traditional x-ray sources.

All three x-ray beams will be collimated toward the 200mm FOV at isocenter by three collimators that are mounted on the gantry between the sources and detectors. The collimated beams can reduce radiation exposure to regions outside the FOV. The 200mm FOV is much smaller compared to the 500mm FOV in most commercially available CT scanners, but it should be enough to cover many important organs such as hearts within human bodies. Smaller FOV also allows a smaller detector size ( $\sim$ 60% reduction). Smaller detector sizes make it possible to mount three imaging chains on a conventional CT gantry ( $\sim$ 1m in diameter) without overlapping. As a result, three projections can be acquired from the three imaging chains simultaneously.

# **III. NUMERICAL ANALYSES**

# A. SCAN ANGLE

The three identical source arrays are symmetrically positioned around the FOV (see Fig. 2). To acquire a complete



FIGURE 2. Illustration for scan angle analysis of the three distributed x-ray source (red color). The three source arrays are arranged symmetrically around the FOV, which is shown as the inner circular region (grey color).

projection dataset for the FOV, the effective scan angle should be greater than  $\pi + \alpha$  where  $\alpha$  is the fan angle and indicated by the angle  $\angle S_1 OS_4$ . Suppose each source array can cover an angle  $\varphi$ ,

$$\angle S_1 O S_2 = \angle S_3 O S_4 = \angle S_5 O S_6 = \varphi. \tag{1}$$

Then we have

$$\angle S_2 O S_3 = \angle S_4 O S_5 = \angle S_6 O S_1 = (2\pi - 3\varphi)/3.$$
 (2)

For a short scan (i.e.  $\pi + \alpha$ ) from  $S_1$  to  $S_4$ , the data from  $S_2$  to  $S_3$  are missing. However, the missing data can be compensated by the source array from  $S_5$  to  $S_6$ . Therefore, we have

$$\begin{cases} \angle S_1 O S_4 = \pi + \alpha \\ \angle S_1 O S_4 = \angle S_1 O S_2 + \angle S_2 O S_3 + \angle S_3 O S_4 \\ = \varphi + (2\pi - 3\varphi)/3 + \varphi. \end{cases}$$
(3)

We immediately arrive at

$$\varphi = \pi/3 + \alpha. \tag{4}$$

As a result, to satisfy the data requirement for a short scan, each source array should cover an angle of  $\pi/3+\alpha$ , and the corresponding detector need to be rotated an angle of  $\pi/3+\alpha$ . From the key geometry parameters listed in Table 1, the fan angle  $\alpha$  can be determined to be 21°.

### **B. TEMPORAL RESOLUTION**

Based on the above analysis in subsection III.1, the temporal resolution can be easily determined for a short scan in the SSRDCT architecture as  $(\pi/3+\alpha)/2\pi * T_{rot}$ , where  $T_{rot}$  is the gantry rotation time. In contrast, the temporal resolutions for short scan of the DSCT and the conventional single-source CT (SSCT) architectures are  $(\pi/2+\alpha)/2\pi * T_{rot}$ , and  $(\pi + \alpha)/2\pi * T_{rot}$  respectively. For half scan (i.e. 180° scan), the temporal resolution would be  $T_{rot}/6$ ,  $T_{rot}/4$ , and  $T_{rot}/2$  for SSRDCT, DSCT, and SSCT, respectively. The relative

# TABLE 2. Comparison of relative temporal resolutions for the SSCT, DSCT, and STRICT architectures. The gantry rotation time was assumed to be same and the SID is 540 mm.

Architecture	SSCT	DSCT	SSRDCT
Number of Source-Detector Chains	1	2	3
Field of View (mm)	500	340 <sup>a</sup>	200
Relative Temporal Resolution – Short Scan <sup>b</sup>	1	0.54	0.35
Relative Temporal Resolution – Half Scan <sup>b</sup>	1	0.50	0.33

<sup>a</sup> The two detectors in DSCT have different sizes and the system FOV is limited by the smaller

detector at 340mm [4].

<sup>b</sup> Temporal resolutions are normalized to those in the traditional SSCT architecture.

temporal resolutions for the three architectures are listed in Table 2, where the gantry rotation time  $T_{rot}$  was assumed to be same and the SID was taken as 540mm. Compared to the latest DSCT architecture, the temporal resolution of SSRDCT architecture is improved by a factor of more than 1/3 for both the short-scan and half-scan modes. If the gantry rotation time  $T_{rot}$  is assumed as 270ms, the temporal resolution of the SSRDCT architecture will be 61ms for the short-scan mode and 45ms for the half-scan mode.

# C. RADIATION DOSE

At the same SID, the radiation dose at isocenter is proportional to the total radiation exposure during a CT scan. For a same SID, a constant x-ray flux can be assumed at the isocenter for a given x-ray source technology. Therefore, the radiation dose at isocenter is solely dependent on the total exposure time. For the SSCT architecture, the x-ray source is turned on continuously during the data acquisition for a single slice image, thus the total exposure time per CT scan equals to the corresponding temporal resolution. For the DSCT architecture, since the two sources are turned on continuously during the data acquisition, the total exposure time is two times of its temporal resolution. For the SSRDCT architecture, each x-ray source will give a single pulsed exposure, thus the total exposure time is the summation of x-ray exposure times from individual x-ray sources in the three source arrays. This total exposure time is not necessarily three times of its temporal resolution, because an 'off time' (i.e. a time period without exposure) can be introduced between two consecutive pulsed exposures from two neighboring x-ray sources. The off time will significantly reduce the total exposure time per CT scan in the SSRDCT architecture. This can help save the radiation dose.

To reduce radiation exposure to patients, many efforts have been devoted to develop image reconstruction algorithms for fewer projection views, less of a scanning arc or lower x-ray intensity per view [26], [31], [32]. These algorithms are highly synergistic to our proposed architecture. They would permit a decent image quality to be obtained in the SSRDCT architecture with less x-ray sources in each source array, smaller coverage angle per array, and lower x-ray source power, resulting in a CT system that is more compact and at lower cost. Particularly, lowering x-ray intensity per view will enable the stationary distributed x-ray sources to adopt the fixed-anode design, which will dramatically simplify the source design and reduce the demand on the amount of electron currents from the cathodes. In the following section, we will evaluate the performance of the architecture with a clinically acquired cardiac CT dataset and the popular total variation minimization (TVM) method combined with steepest descent (TVM-SD) search [33], [34].

# **IV. SIMULATION AND RESULTS**

To evaluate the image quality of proposed CT architecture, we carried out numerical simulations with clinically acquired cardiac CT imaging data in a typical fan-beam geometry. The data were acquired on a commercial CT scanner (GE Discovery CT750 HD) at Wake Forest University Health Science. The radius of the scanning trajectory is 538.5mm. 2200 projections were uniformly acquired over a full scan of 360°. Each projection has 888 equiangular distributed detector elements. The popular total variation minimization (TVM) with steepest descent (TVM-SD) search algorithm [33], [34] was used to reconstruct a global image (Figure 3 (a)), which serves as a benchmark to evaluate the image quality of the SSRDCT architecture. The global



FIGURE 3. Reconstructed slice images using the TVM-SD algorithm with a display window [-1000HU, 1800HU]. (a) Is the reconstructed global image from full dataset after 60 iterations, (b) and (c) are the reconstructed ROI images for the short-scan mode and half-scan mode in the SSRD-CT architecture after 200 iterations.

image is a matrix of  $512 \times 512$  pixels and covers a region of  $498.3 \times 498.3$  mm<sup>2</sup>.

To simulate an interior-ROI-oriented data acquisition, each projection was truncated by discarding 270 detector elements from each of the two sides. This resulted in an interior ROI of 200mm in diameter with a fan angle of 21.4°. Therefore, for a short scan the cover angle of each source array should be 81.4°, and only the acquired projections in  $[0, 81.4^{\circ}]$ , [120°, 201.4°], and [240°, 321.4°] are needed in the SSRDCT architecture. Correspondingly, for a half scan only the projections in [0°, 60°], [120°, 180°], and [240°, 300°] are needed. These selected projections were first truncated, and then used to reconstruct interior images using the TVM-SD algorithm. Figure 3(b) and 3(c) show the reconstructed ROI images for the short-scan and half-scan modes, respectively. Figure 4 shows the representative line profiles along the horizontal and vertical central lines inside the ROI in the three images in Figure 3. We can see there is an excellent match between the global image and the ROI images.

The TVM-SD algorithm was developed based on the CS theory. According to the CS theory, high-quality images can be reconstructed from a limited number of projections. This means fewer views can be used (thus reducing radiation dose). For the half-scan mode (i.e. 60° coverage angle per source array), we down-sampled the projections using the global benchmark image. The projections views for the x-ray sources in the three source arrays were equiangularly spaced in the [0, 60°], [120°, 180°] and [240°, 300°] angular ranges. The number of views per source array was set to be 160, 130, 100, 70 and 40, respectively, and the corresponding reconstructed ROI images are shown in Figure 5. Figure 6 shows two representative line profiles along horizontal and vertical central lines. From Figures 5 and 6, we can see that the reconstructed images from 160, 130 and 100 views per array are all qualitatively comparable with the benchmark, while the others lose some details as indicated by the arrows in Figure 5 (d) and (e).

To quantitatively evaluate the quality of reconstructed images from the half-scan mode, we calculated the rootmean-square error (RMSE) in the ROI region between the reconstructed ROI images and the benchmark. The results are shown in Figure 7. We can see that when the iteration number reaches 1000, the RMSEs of  $160 \times 3$ ,  $130 \times 3$  and  $100 \times 3$  views are significantly smaller than those of  $70 \times 3$  and  $40 \times 3$  views. Its implication for dose reduction is highly significant. If we use the result from  $100 \times 3$  views as a reference, this means that the radiation dose could be potentially reduced by a factor of 3.7.

### **V. DISCUSSION AND CONCLUSION**

In this paper, a SSRDCT architecture is presented by combining two latest advances in the CT field: the interior tomography [26] and the distributed x-ray source [27], [28]. The feasibility is investigated via extensive numerical analyses and simulations using clinical cardiac CT data. Compared to the current  $3^{rd}$ -generation CT architecture, the SSRDCT has a great potential to reach  $\leq$ 50ms temporal resolution



FIGURE 4. Representative line profiles of the images in Figure 3 along the horizontal and vertical directions at the center.



**FIGURE 5.** Reconstructed ROI images for the half-scan mode with few views in a display window [-1000HU, 1800HU]. The Iteration numbers for all the images are 1000, and the total number of views are 160 × 3, 130 × 3, 100 × 3, 70 × 3, and 40 × 3 for (a)–(e), respectively.

and reduce radiation dose significantly. The enhanced performance metrics have high diagnosis values for clinical exams especially for cardiovascular diseases (CVDs). Indeed, cardiac CT has been used to help diagnose a range of CVDs including coronary heart disease, valvular heart disease, pericardial diseases, congenital heart disease, and pulmonary vein disease [35].

Previously, high CT temporal resolution has been sought out by two approaches - DSR and EBCT. While fast temporal resolution in the DSR was achieved by increasing the number of source-detector chains, for the EBCT, fast temporal resolution was obtained through a stationary x-ray source. The fast temporal resolution in SSRDCT can be explained from the following three points. First, it is well known that simultaneous data acquisition from multiple source-detector chains can boost temporal resolution. The SSRDCT has three source-detector chains, which are expected to boost temporal resolution in a fashion similar to the DSR [15]–[17]. Second, the three distributed x-ray sources in SSRDCT are stationary, and scanning x-ray beams are realized electronically rather mechanically. The stationary sources will deliver the same benefit as that shown in the EBCT [13]. Compared to mechanical rotation, electronic sweeping allows the source-point to be moved at a far greater speed. Lastly, by removing the vacuum-based x-ray sources and their power electronics away from the mechanical gantry, what are left on the gantry will be the semiconductor-based detectors and their low-power electronics. The engineering challenge to spin a heavily loaded gantry will be gone. This design will not only reduce the mechanical complexity for the gantry but also could lead to a much faster gantry rotation speed (i.e. smaller  $T_{rot}$ ) and hence an even higher temporal resolution in SSRDCT.

Historically, the high temporal resolution in the DSR and EBCT was not achieved without some drawbacks. Because the DSR had to employ a huge gantry to mount multiple source-detector chains, it came with large size and high cost, and only one such system has been ever built. The EBCT has an anode ring that encloses the patient body. To steer an electron beam to strike on the large target ring, the EBCT system also came with big size and high cost. The proposed SSRDCT is based on a conventional CT gantry, due to the stationary distributed x-ray source and the smaller detectors. The resulted system will be compact and will have a size similar to the current CT systems. As a result, the SSRDCT can achieve a high temporal resolution without the aforementioned drawbacks of DSR and EBCT. Additionally, the EBCT is also known for its other drawbacks from a stationary fullring detector. A stationary full-ring detector, in addition to its high cost, also requires an offset in the z-direction between the source and detector planes to cover a typical scan angle of 220°. This configuration does not allow a typical antiscatter grid collimator on the detector. In contrast, because the x-ray beams and the detectors are co-planar in the SSRDCT, detector anti-scatter grids can be still used. This is made possible because the rotating detectors are always at the opposite sides of the sweeping source-points. The sweeping source points can be realized by combining the electronic switching



FIGURE 6. Two representative profiles along horizontal and vertical central lines for the images in Figure 5.



FIGURE 7. RMSEs vs. iteration numbers for different numbers of views at the half-scan mode.

of the gate voltages on the multiple cathodes and the local steering (i.e. steering electronically and/or magnetically the electron stream from a cathode to strike onto short focal track on the fixed anode during the exposure time for a projection).

The dose reduction arises from the interior scan and the CS-based interior reconstruction algorithm. The smaller an ROI, the narrower the beam, and thus less radiation. Narrower x-ray beams also reduce photon scattering to improve contrast resolution. The fewer views a CT scan acquires, the smaller number of x-ray beam assemblies will be required in a distributed x-ray source, and hence lower construction cost for the source. The distributed x-ray sources allow swift pulsing of the x-ray beams, which can eliminate the unnecessary x-ray exposures to save radiation dose. Therefore, the state-of-the-art few-view based interior tomography is highly synergistic with the distributed x-ray sources technology. To realize the benefit in lower radiation dose, we need to collimate the scanning x-ray beams from the stationary distributed x-ray sources during a CT scan. This can be realized by mounting three x-ray beam collimators on the rotating gantry and opposite to the three detectors. This arrangement can ensure that an x-ray beam emitted from the active source will be always well aligned with the corresponding collimator and detector.

The proposed SSRDCT architecture is not perfect and it has the following limitations. First, it has a small FOV (200mm), which is not sufficient to cover a full width of human body. Second, the critical components - the distributed x-ray sources – pose some engineering challenges. For the first limitation, the hospitals may need a dedicated heartcentered CT architecture because of the prevalence and morbidity of CVDs. The potential in higher temporal resolution and lower radiation dose is especially attractive for cardiac CT perfusion imaging studies. A heart-dedicated CT scanner, even though it can only cover 200mm FOV, can also find it suitable to evaluate other vital organs such as kidneys and livers. For the second limitation, while such sources are not available yet, recent progresses in CNT-based distributed x-ray sources provided a high possibility for their implementations. A linear x-ray tube with 75 CNT-based x-ray beams has been constructed to test some other interesting CT configurations [36]. Another linear x-ray tube with 31 individually controllable x-ray sources has been developed for stationary digital breast tomosynthesis [29]. It is our hope that the potential improvements in speed and dose of this SSRDCT architecture will promote the realization of such x-ray sources sooner.

Finally, other variants (in addition to the specific type in Figure 1) can be designed in the general frame of the SSRDCT architecture. An obvious extension is to use a stationary ring source that covers a full circle. This will make it possible for a full-scan. However, it will have higher cost on the source, require each detector to rotate at least 120°, and hence lower the temporal resolution. Another possibility is to use linear source arrays to replace curved source arrays. This design will lead to geometrical nonuniformity from view to view, but it could lower the cost for the distributed x-ray sources. Designing a new CT architecture is a complex task. At this stage, we mainly focused on identifying some feasible sampling geometry (coverage angles and view numbers etc.) and the corresponding image reconstruction algorithms. Some practical issues could have been not adequately investigated. For example, there may not be enough gap distance between the sources and detectors.

In conclusion, we present and evaluate a new SSRDCT architecture that is expected to provide higher temporal resolution and lower radiation dose. The SSRDCT architecture synergistically combines the few-view interior reconstruction algorithms and the distributed x-ray source technology. Although there may be some engineering challenges for its implementation, based on our analysis and simulation, the proposed SSRDCT architecture has a high potential to outperform the current CT architecture in both temporal resolution and radiation dose. According to the World Health Organization (WHO), the number of deaths from CVDs is estimated to be about 20 million in 2015 [37]. The heart-dedicated CT scanners based on the proposed SSRDCT architecture could be useful for our global mission to reduce deaths from CVDs.

# REFERENCES

- G. N. Hounsfield, "Computerized transverse axial scanning (tomography): Part 1. Description of system," *Brit. J. Radiol.*, vol. 46, no. 552, pp. 1016–1022, Feb. 1973.
- [2] W. A. Kalender, Computed Tomography: Fundamentals, System Technology, Image Quality, Applications, 2nd ed. Erlangen, Germany: Publicis Corporate Pub. 2005.
- [3] M. Lell et al., "Prospectively ECG-triggered high-pitch spiral acquisition for coronary CT angiography using dual source CT: Technique and initial experience," Eur. Radiol., vol. 19, no. 11, pp. 2576–2583, 2009.
- [4] T. G. Flohr *et al.*, "Dual-source spiral CT with pitch up to 3.2 and 75 ms temporal resolution: Image reconstruction and assessment of image quality," *Med. Phys.*, vol. 36, no. 12, pp. 5641–5653, 2009.
- [5] S. Achenbach *et al.*, "Contrast-enhanced coronary artery visualization by dual-source computed tomography—Initial experience," *Eur. J. Radiol.*, vol. 57, no. 3, pp. 331–335, 2006.
- [6] T. G. Flohr *et al.*, "First performance evaluation of a dual-source CT (DSCT) system," *Eur. Radiol.*, vol. 16, no. 2, pp. 256–268, 2006.
- [7] T. R. Johnson *et al.*, "Dual-source CT cardiac imaging: Initial experience," *Eur. Radiol.*, vol. 16, no. 7, pp. 1409–1415, May 2006.
- [8] M. Kachelrie, M. Knaup, and W. A. Kalender, "Multithreaded cardiac CT," Med. Phys., vol. 33, no. 7, pp. 2435–2447, 2006.

- [9] H. Scheffel *et al.*, "Accuracy of dual-source CT coronary angiography: First experience in a high pre-test probability population without heart rate control," *Eur. Radiol.*, vol. 16, no. 12, pp. 2739–2747, 2006.
- [10] S. Leschka et al., "Noninvasive coronary angiography with 64-section CT: Effect of average heart rate and heart rate variability on image quality," *Radiology*, vol. 241, no. 2, pp. 378–385, Sep. 2006.
- [11] M. Mahesh and D. D. Cody, "Physics of cardiac imaging with multiplerow detector CT," *RadioGraphics*, vol. 27, no. 5, pp. 1495–1509, Sep./Oct. 2007.
- [12] P. Schardt *et al.*, "New X-ray tube performance in computed tomography by introducing the rotating envelope tube technology," *Med. Phys.*, vol. 31, no. 9, pp. 2699–2706, 2004.
- [13] M. J. Lipton, C. B. Higgins, D. Farmer, and D. P. Boyd, "Cardiac imaging with a high-speed cine-CT scanner: Preliminary results," *Radiology*, vol. 152, no. 3, pp. 579–582, Sep. 1984.
- [14] W. A. Kalender, "CT: The unexpected evolution of an imaging modality," *Eur. Radiol.*, vol. 15, pp. D21–D24, Nov. 2005.
- [15] E. L. Ritman, J. H. Kinsey, R. A. Robb, B. K. Gilbert, L. D. Harris, and E. H. Wood, "Three-dimensional imaging of heart, lungs, and circulation," *Science*, vol. 210, no. 4467, pp. 273–280, 1980.
- [16] E. L. Ritman, L. D. Harris, J. H. Kinsey, and R. A. Robb, "Computed tomographic imaging of the heart: The dynamic spatial reconstructor," *Radiol. Clin. North Amer.*, vol. 18, no. 3, pp. 547–555, Dec. 1980.
- [17] E. L. Ritman, R. A. Robb, and L. D. Harris, *Imaging Physiological Func*tions: Experience With the DSR. Philadelphia, PA, USA: Praeger, 1985,
- [18] P. S. Douglas *et al.*, "Developing an action plan for patient radiation safety in adult cardiovascular medicine: Proceedings from the Duke University clinical research Institute/American college of cardiology foundation/American heart association think tank held on February 28, 2011," *J. Amer. College Cardiol.*, vol. 59, no. 20, pp. 1833–1847, 2012.
- [19] R. P. Marcus, K. Nikolaou, D. Theisen, M. F. Reiser, and F. Bamberg, "Myocardial perfusion imaging by computed tomography: Today and tomorrow," *Int. J. Clin. Pract. Suppl.*, vol. 65, no. 173, pp. 14–22, 2011.
- [20] T. G. Flohr, R. Raupach, and H. Bruder, "Cardiac CT: How much can temporal resolution, spatial resolution, and volume coverage be improved?" *J. Cardiovascular Comput. Tomography*, vol. 3, no. 3, pp. 143–152, May/Jun. 2009.
- [21] R. E. van Gelder *et al.*, "CT colonography: Feasibility of substantial dose reduction—Comparison of medium to very low doses in identical patients," *Radiology*, vol. 232, no. 2, pp. 611–620, 2004.
- [22] H. Yu, S. Zhao, E. A. Hoffman, and G. Wang, "Ultra-low dose lung CT perfusion regularized by a previous scan," *Acad. Radiol.*, vol. 16, no. 3, pp. 363–373, 2009.
- [23] J. E. Tkaczyk, Y. Du, D. J. Walter, X. Wu, J. Li, and T. Toth, "Simulation of CT dose and contrast-to-noise as function of bowtie shape," *Proc. SPIE*, vol. 5368, pp. 403–410, May 2004.
- [24] C. Suess and X. Chen, "Dose optimization in pediatric CT: Current technology and future innovations," *Pediatric Radiol.*, vol. 32, no. 10, pp. 729–734, 2002.
- [25] P. J. La Rivière, "Penalized-likelihood sinogram smoothing for lowdose Ct," *Med. Phys.*, vol. 32, no. 6, pp. 1676–1683, 2005.
- [26] H. Yu and G. Wang, "Compressed sensing based interior tomography," *Phys. Med. Biol.*, vol. 54, no. 9, pp. 2791–2805, 2009.
- [27] K. Frutschy et al., "High power distributed X-ray source," Proc. SPIE, vol. 7622, p. 76221H, Mar. 2010.
- [28] F. Sprenger et al., "Stationary digital breast tomosynthesis with distributed field emission X-ray tube," Proc. SPIE, vol. 7961, p. 878280, Mar. 2011.
- [29] X. Qian *et al.*, "High resolution stationary digital breast tomosynthesis using distributed carbon nanotube X-ray source array," *Med. Phys.*, vol. 39, no. 4, pp. 2090–2099, 2012.
- [30] J. Zhang, G. Yang, Y. Z. Lee, S. Chang, J. P. Lu, and O. Zhou, "Multiplexing radiography using a carbon nanotube based X-ray source," *Appl. Phys. Lett.*, vol. 89, no. 6, p. 064106, 2006.
- [31] E. Y. Sidky and X. Pan, "Image reconstruction in circular cone-beam computed tomography by constrained, total-variation minimization," *Phys. Med. Biol.*, vol. 53, no. 17, pp. 4777–4807, 2008.
- [32] G.-H. Chen, J. Tang, B. Nett, Z. Qi, S. Leng, and T. Szczykutowicz, "Prior image constrained compressed sensing (PICCS) and applications in X-ray computed tomography," *Current Med. Imag. Rev.*, vol. 6, no. 2, pp. 119–134, May 2010.
- [33] H. Yu and G. Wang, "A soft-threshold filtering approach for reconstruction from a limited number of projections," *Phys. Med. Biol.*, vol. 55, no. 3, pp. 3905–3916, 2010.

- [34] B. Liu et al., "Image reconstruction from limited angle projections collected by multisource interior X-ray imaging systems," *Phys. Med. Biol.*, vol. 56, no. 19, pp. 6337–6357, 2011.
- [35] M. Y. Desai, "Cardiac CT beyond coronary angiography: Current and emerging non-coronary cardiac applications," *Heart*, vol. 97, no. 3, pp. 417–424, 2011.
- [36] T. Zhang, D. Schulze, X. Xu, and J. Kim, "Tetrahedron beam computed tomography (TBCT): A new design of volumetric CT system," *Phys. Med. Biol.*, vol. 54, no. 11, pp. 3365–3378, 2009.
- [37] WHO. (2005). Preventing Chronic Diseases: A Vital Investment. [Online]. Available: http://www.who.int/chp/chronic\_disease\_report/ full\_report.pdf



**GUOHUA CAO** is currently an Assistant Professor of Biomedical Engineering with the Virginia Tech-Wake Forest University School of Biomedical Engineering and Sciences, Virginia Polytechnic Institute and State University (Virginia Tech), Blacksburg, VA, USA. He received the Ph.D. degree in physical chemistry from Brown University, Providence, RI, USA, in 2005. From 2005 to 2008, he pursued two post-doctoral trainings at Brown University and the University of

North Carolina at Chapel Hill (UNC-Chapel Hill), Chapel Hill, NC, USA, in research related to X-ray imaging. From 2009 to 2011, he was a Research Assistant Professor with the Department of Physics, UNC-Chapel Hill. He joined Virginia Tech as a tenure-track faculty in 2011. He has received a number of awards, including the UNC's Post-Doctoral Scholar Award for Research Excellence, the Outstanding Poster Award from the National Cancer Institute Alliance for Nanotechnology in Cancer Investigators Meeting in 2007, and the Honorable Mention Poster Award from the International Society for Optical Engineering - Medical Imaging Conference in 2009. He was also a recipient of the NSF CAREER Award in 2014. His research is directed at medical imaging, with a focus on developing novel imaging systems and methods based on carbon nanotube field-emission X-ray technology.



**BAODONG LIU** is currently an Associate Professor with the Key Laboratory of Nuclear Analysis Techniques, Institute of High Energy Physics, Chinese Academy of Sciences, Beijing, China, and the Beijing Engineering Research Center of Radiographic Techniques and Equipment, Beijing. He received the Ph.D. and bachelor's degrees from Chongqing University, Chongqing, China, in 2011, 2008, and 2005, respectively. From 2011 to 2013, he was a Post-

Doctoral Research Fellow with the Division of Biomedical Imaging, School of Biomedical Engineering and Sciences, Wake Forest University, Winston-Salem, NC, USA.



**HAO GONG** is currently pursuing the Ph.D. degree with the School of Biomedical Engineering and Sciences, Virginia Polytechnic Institute and State University, Blacksburg, VA, USA. He received the master's degree in electrical and computer engineering from Purdue University, West Lafayette, IN, USA, in 2010, and the bachelor's degree from the School of Information Science and Engineering, Central South University, Changsha, China, in 2007. From 2011 to

2012, he was a Research Engineer with the Department of Electrical and Computer Engineering, Purdue University. In 2012, he was selected as a Pratt Scholar at the Virginia Polytechnic Institute and State University. His current research interests include X-ray computed tomography and medical image processing with an emphasis on system architecture design of X-ray computed tomography.



**HENGYONG YU** (SM'06) is currently an Assistant Professor and the Director of the CT Laboratory with the Department of Biomedical Engineering, Wake Forest University Health Sciences, Winston-Salem, NC, USA. He received the bachelor's degrees in information science and technology, and computational mathematics, and the Ph.D. degree in information and communication engineering from Xi'an Jiaotong University, Xi'an, China, in 1998, 1998, and 2003, respec-

tively. He was an Instructor and Associate Professor with the College of Communication Engineering, Hangzhou Dianzi University, Hangzhou, China, from 2003 to 2004. From 2004 to 2006, he was a Post-Doctoral Fellow and an Associate Research Scientist with the Department of Radiology, University of Iowa, Iowa City, IA, USA. From 2006 to 2010, he was a Research Scientist and the Associate Director of CT Laboratory, Division of Biomedical Imaging, Virginia Tech-Wake Forest University School of Biomedical Engineering and Sciences, Virginia Polytechnic Institute and State University, Blacksburg, VA, USA. He joined Wake Forest University Health Sciences as a faculty member in 2010. His interests include computed tomography and medical image processing. He has authored or co-authored over 100 peer-reviewed journal papers with an H-index of 19 according to the web of knowledge. He is the founding Editor-in-Chief of JSM Biomedical Imaging Data Papers, serves as an Editorial Board Member of Signal Processing, the Journal of Medical Engineering, CT Theory and Applications, the International Journal of Biomedical Engineering and Consumer Health Informatics, and Open Medical Imaging Journal, and the Guest Editor of the IEEE TRANSACTIONS ON MEDICAL IMAGING, the IEEE Access, and the International Journal of Biomedical Imaging. He is a Senior Member of the IEEE Engineering in Medicine and Biology Society, and a member of the American Association of Physicists in Medicine and the Biomedical Engineering Society. He was a recipient of the Outstanding Doctoral Dissertation Award from Xi'an Jiaotong University, Xi'an, China, in 2005, the Best Natural Science Paper Award from the Association of Science and Technology of Zhejiang Province, and the NSF CAREER Award for development of CS-based interior tomography in 2012.



**GE WANG** (F'03) received the Ph.D. degree in electrical, computer, and systems engineering from the Rensselaer Polytechnic Institute, Troy, NY, USA, where he is currently a Clark and Crossan Endowed Chair Professor and the Director of the Biomedical Imaging Center/Cluster. His expertise includes X-ray computed tomography (CT) and optical molecular tomography. He wrote the pioneering papers on the first spiral cone-beam CT algorithm (1991 and 1993) that enables spi-

ral/helical cone-beam CT imaging, which is constantly used in almost all hospitals worldwide. There are over 70-million CT scans yearly in the U.S. alone, with a majority in the spiral cone-beam/multislice mode. He and his collaborators also wrote the first paper on bioluminescence tomography, creating a new area of optical molecular tomography. His group published the first papers on interior tomography and omnitomography for grand fusion of all relevant tomographic modalities (all-in-one) to acquire different datasets simultaneously (all-at-once) and capture multiphysics interactions (all-ofcouplings) with simultaneous CT-MRI and simultaneous CT-SPECT as special examples. His results were featured in Nature, Science, and Proceedings of the National Academy of Sciences, and recognized with various academic awards. He has authored about 400 journal papers, which are highly cited. His group has been in close collaboration with multiple world-class groups, and continuously funded by federal agents (25MasPI/ContactPI/Multi - PI, 28M as Co-PI/Co-I/Mentor). He is the lead Guest Editor of four IEEE TRANSACTIONS ON MEDICAL IMAGING special issues on X-ray CT, molecular imaging, compressive sensing, and spectral CT, respectively, the founding Editor-in-Chief of the International Journal of Biomedical Imaging, an Associate Editor of the IEEE TRANSACTIONS ON MEDICAL IMAGING, Medical Physics Journal, and others. He is a fellow of the International Society for Optical Engineering, the Optical Society of Amercica, the American Institute for Medical and Biological Engineering, and the American Association of Physicists in Medicine.