

First in vivo head-to-head comparison of high-definition versus standard-definition stent imaging with 64-slice computed tomography

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Abstract The aim of this study was to compare image quality characteristics from 64-slice high definition (HDCT) versus 64-slice standard definition CT (SDCT) for coronary stent imaging. In twenty-five stents of 14 patients, undergoing contrast-enhanced CCTA both on 64-slice SDCT (LightSpeedVCT, GE Healthcare) and HDCT (Discovery HD750, GE Healthcare), radiation dose, contrast, noise and stent characteristics were assessed. Two blinded observers graded stent image quality (score 1 = no, 2 = mild, 3 = moderate, and 4 = severe artefacts). All scans were reconstructed with increasing contributions of adaptive statistical iterative reconstruction (ASIR) blending (0, 20, 40, 60, 80 and 100 %). Image quality was significantly superior in HDCT versus SDCT (score 1.7 ± 0.5 vs. 2.7 ± 0.7 ; $p < 0.05$). Image noise was significantly higher in HDCT compared to SDCT irrespective of ASIR contributions ($p < 0.05$). Addition of 40 % ASIR or more reduced image noise significantly in both HDCT and SDCT. In HDCT in-stent luminal attenuation was significantly lower and mean measured in-stent luminal diameter was significantly larger (1.2 ± 0.4 mm vs. 0.8 ± 0.4 mm; $p < 0.05$) compared to SDCT. Radiation dose from HDCT was comparable to SDCT (1.8 ± 0.7 mSv vs. 1.7 ± 0.7 mSv; $p = \text{ns}$). Use of

HDCT for coronary stent imaging reduces partial volume artefacts from stents yielding improved image quality versus SDCT at a comparable radiation dose.

Keywords Coronary computed tomography angiography · Stent imaging · High definition CT · Iterative reconstruction

Introduction

Coronary computed tomography angiography (CCTA) has become a well-established non-invasive tool for diagnosis of coronary artery disease (CAD). While yielding high accuracy compared to invasive coronary angiography for the detection of coronary artery stenosis [1, 2], CCTA for stent imaging is still affected by artefacts from coronary stents leading to artificial luminal narrowing, due to partial volume artefact from highly attenuated stent struts [3, 4].

Although the in-stent restenosis rate has been substantially reduced by introducing drug-eluting stents it has not been entirely eliminated [5]. Furthermore, there is an increasing population of stented patients who are evaluated for progression of CAD. Therefore, any refinements in low radiation dose CCTA [2, 6] which may offer improved assessment of stents would be welcome.

Recently a new generation of high definition CT (HDCT) scanner (Discovery CT 750 HD, GE Healthcare, Milwaukee) has been introduced with substantially improved in plane resolution ($0.23 \text{ mm} \times 0.23 \text{ mm}$) complemented by a new adaptive statistical iterative reconstruction (ASIR, GE Healthcare) algorithm in order to compensate for the increased noise due to the higher resolution [7]. Preliminary in vivo results have suggested that HDCT may provide superior evaluation of coronary stents

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compared with SDCT [8]. Recent in vivo results seem to confirm that HDCT may improve image quality for coronary stents with 2.75–3.5 mm diameter compared with conventional standard definition CT (SDCT) due to higher spatial resolution [9], but those results were obtained from two different patient populations, one examined on a HDCT and the other on SDCT. Furthermore, the HDCT scans were reconstructed with latest reconstruction algorithms such as ASIR, while SDCT images were not. Thus, from that study, with relatively high radiation dose due to retrospective triggering, no firm conclusion on stent image quality from HDCT can be drawn. A recent elegant study has compared prospectively triggered low dose SDCT and HDCT using invasive coronary angiography and intravascular ultrasound as gold standard but did not provide head-to-head comparison of the two scanners in the same patients [10].

Therefore, the purpose of the present study was to compare the performance of 64-slice HDCT to 64-slice SDCT for coronary stent imaging of the same patients by assessing image quality as well as quantitative parameters, such as stent geometry and in-stent contrast attenuation.

Materials and methods

Study population

The study protocol was approved by the ethics committee, and written informed consent was obtained from all patients. Fourteen patients, who were referred by clinical indication for the assessment of CAD and had undergone previous percutaneous coronary intervention (PCI) with stent implantation were prospectively enrolled in the study. The patients underwent prospectively triggered contrast-enhanced CCTA both on a 64-slice SDCT (LightSpeed VCT, GE Healthcare) and HDCT scanner (Discovery HD 750, GE Healthcare) on the same day. Exclusion criteria were known hypersensitivity to iodinated contrast agent, renal insufficiency (glomerular filtration rate < 60 ml/min), non-sinus rhythm.

CCTA acquisition and reconstruction

Prior to the examination metoprolol (up to 25 mg Beloc, Atrazeneca, London, UK) was injected intravenously if heart rate was higher than 65 beats per minute and 2.5 mg isosorbiddinitrate (Isoket, Schwarz Pharma, Monheim, Germany) was administered sublingually in order to obtain optimal image quality for CCTA. All patients underwent contrast-enhanced CCTA during inspiration breath hold with prospective ECG-triggering as previously reported [2, 11, 12] on a SDCT (LightSpeedVCT, GE Healthcare) and a

HDCT (Discovery HD 750, GE Healthcare) scanner. Iodixanol (Visipaque 320, 320 mg/ml, GE Healthcare) was injected into an antecubital vein followed by 50 ml saline solution. Contrast media volume (40–105 ml) and flow rate (3.5–5 ml/s) were adapted to body surface area (BSA) as previously validated [13].

Scanning parameters of SDCT and HDCT were as follows: Collimation of 64×0.625 mm, gantry rotation time of 0.35 s and field of view was 25 cm. Tube voltage (100–120 kV) and tube current (450–700 mA) were adapted to body mass index as previously described [14]. Scans of HDCT were acquired in high resolution mode with an in-plane spatial resolution of $0.23 \text{ mm} \times 0.23 \text{ mm}$ as previously reported [7]. All scans of SDCT and HDCT were reconstructed using FBP (0 % ASIR) and increasing ASIR blending factors, i.e. 20, 40, 60, 80 and 100 %. The iterative reconstruction algorithm ASIR has been described in detail previously [15–17]. In brief, ASIR reconstructs pictures by comparing measured projection with a synthesized projection using both statistical fluctuation calculations and system optics.

Radiation dose

Effective radiation dose from CCTA was calculated as the product of dose-length product (DLP) times a conversion coefficient for chest ($k = 0.014 \text{ mSv mGy}^{-1} \text{ cm}^{-1}$) [18].

Stent visualization and image quality

All images were transferred to an external workstation (AW 4.6, GE Healthcare) for analysis. Qualitative stent analysis was performed by two independent, experienced coronary CCTA readers, blinded for scanner type, clinical history, indication, and stent characteristics. Axial slices as well as curved multiplanar reformations were analysed using a window width of 1200 HU and window level of 240 HU. Image quality was evaluated on a 4-point Likert score for each stent as previously reported (adapted) [19]. Score 1: no artefacts in the surrounding of the stent; score 2: minor artefacts; score 3: major artefacts partially obscuring the stent surrounding; score 4: severe artefacts affecting diagnostic information. Reasons for impaired image quality (stents scored 2–4) were classified as partial volume artefacts, motion artefacts, or calcifications.

Attenuation measurements

Image signal and noise were determined by placing a circled region of interest (ROI) with 20 mm diameter in the aortic root and defined as the mean attenuation value and its standard deviation. Furthermore attenuation was measured within the stent lumen as well as 5 mm proximal and 5 mm distal to the stent. ROIs were drawn as large as

artery (LCX; $n = 2$), in the right coronary artery (RCA; $n = 3$), in the posterior descending artery (PDA; $n = 3$) and in side branches ($n = 2$). There were 10 different types of stents implanted in the patients, i.e. Cypher (Cordis, Miami, FL, USA; $n = 8$), Resolute Integrity (Medtronic Vascular, Santa Rosa, CA; $n = 4$), Endeavor (Medtronic, Minn; $n = 3$), Biomatrix (Biosensors Interventional Technologies Pte Ltd., Singapore; $n = 3$), Presillion (Cordis Corp., Miami, FL; $n = 1$), PRO-Kinetic (Biotronic, Switzerland; $n = 1$), Driver (Medtronic, Natick, MA, USA; $n = 2$), Xience prime (Abbott Vascular, Santa Clara, CA, USA; $n = 1$), Nobori (Terumo Corporation, Tokyo, Japan; $n = 1$) and Taxus (Boston Scientific, Boston, MA, USA; $n = 1$).

Radiation dose

The mean radiation dose of CCTA was 1.8 ± 0.7 mSv from HDCT and 1.7 ± 0.7 mSv from SDCT ($p = ns$).

Image quality

All 25 stents could be visualized with both HDCT and SDCT. In HDCT versus SDCT stents without artefacts (score 1) were found in 32 versus 0 % ($n = 8$ vs. 0), minor artefacts (score 2) were found in 64 versus 44 % ($n = 16$ vs. 11), moderate artefacts (score 3) were found in 4 versus 40 % ($n = 1$ vs. 10) and severe artefacts (score 4) were found in 0 versus 16 % ($n = 0$ vs. 4; Fig. 2). Reasons for impaired image quality in HDCT and SDCT were partial volume artefacts in 32 and 56 % ($n = 8$ and 14), motion artefacts in 4 and 12 % ($n = 1$ and 3) and calcifications in 32 and 32 % ($n = 8$ and 8). The use of HDCT increased the number of stents with no or minor artefacts significantly from 44 to 96 % ($p < 0.05$). Mean image quality score was significantly superior in HDCT versus SDCT for 0, 60, and 100 % ASIR (Table 2; $p < 0.05$), revealing highest image quality by HDCT with 60 % ASIR. Interobserver analysis showed good agreement ($\kappa = 0.8$).

Attenuation measurements

There was no significant difference in mean signal of the ascending aorta in HDCT and SDCT (434.6 ± 85.7 and 438.3 ± 86.1 $p = ns$). Addition of different ASIR contributions had no impact on the latter values in both scanners ($p = ns$). Image noise was significantly higher in HDCT compared with SDCT in all ASIR reconstructions ($p < 0.05$). Compared to FBP reconstruction (0 % ASIR), in both HDCT and SDCT, image noise was significantly reduced using 40 % of ASIR or more, up to a noise reduction of 48 % in HDCT and 41 % in SDCT (Fig. 3).

In-stent luminal attenuation was significantly lower in HDCT compared to SDCT in all ASIR reconstructions

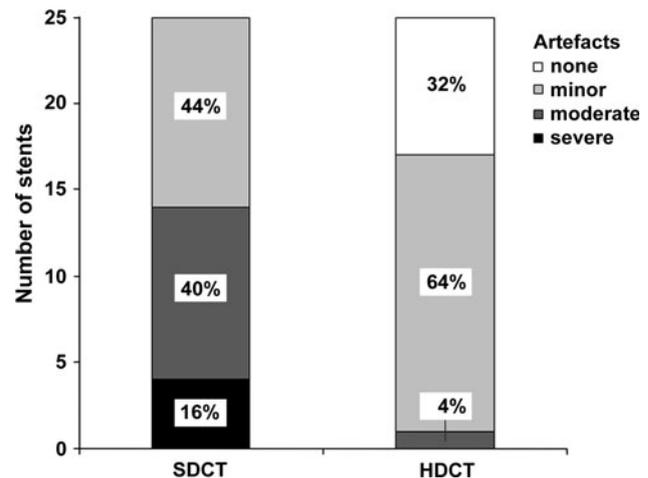


Fig. 2 Image quality score of stents scanned with SDCT and HDCT (0 % ASIR)

Table 2 Comparison of image quality scores in SDCT and HDCT images

Scanner	0 % ASIR	60 % ASIR	100 % ASIR
SDCT	2.7 ± 0.7	2.9 ± 0.6	$3.4 \pm 0.8^+$
HDCT	$1.7 \pm 0.5^*$	$1.5 \pm 0.6^*$	$2.2 \pm 0.8^{*+}$

Image quality scores in standard definition computed tomography (SDCT) and high definition computed tomography (HDCT)

* $p < 0.05$ versus SDCT, + $p < 0.05$ versus 0 % ASIR

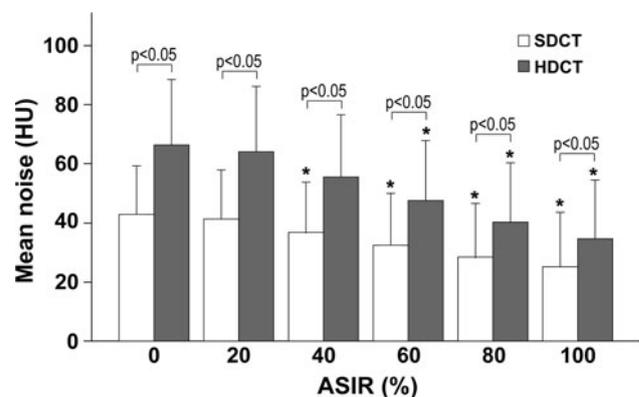


Fig. 3 While there was no significant difference in mean signal of the ascending aorta in SDCT and HDCT, noise was significantly higher in HDCT compared to SDCT ($p < 0.05$). In addition compared to 0 % ASIR noise was significantly reduced by increasing contributions of ASIR (* $p < 0.05$)

($p < 0.05$), whereas no significant difference was measured proximal or distal to the stents (Table 3).

Stent geometry

The mean measured in-stent luminal diameter was significantly larger in HDCT than in SDCT (1.2 ± 0.4 mm vs.

Table 3 Comparison of CT attenuation measurements on SDCT and HDCT images

Location of measurement	Scanner	0 % ASIR	20 % ASIR	40 % ASIR	60 % ASIR	80 % ASIR	100 % ASIR
In-stent lumen	SDCT	598.3 ± 124.2	586.2 ± 126.9	588.8 ± 127.4	591.5 ± 128.1	594.4 ± 129.2	592.0 ± 131.2
	HDCT	474.1 ± 108.1	450.0 ± 110.5	456.0 ± 124.6	453.1 ± 112.2	456.1 ± 110.2	447.7 ± 116.3
		$p < 0.05$					
Proximal to stent*	SDCT	396.0 ± 90.0*	387.1 ± 89.7*	383.9 ± 90.2*	380.7 ± 90.8*	377.6 ± 91.6*	374.3 ± 92.6*
	HDCT	396.1 ± 88.9*	403.4 ± 87.4*	395.5 ± 88*	390.9 ± 87.1*	386.2 ± 83.8*	381.2 ± 83.8*
		$p = ns$					
Distal to stent*	SDCT	351.8 ± 88.8*	360.4 ± 93.3*	351.1 ± 94.6*	342.0 ± 96.4*	332.8 ± 98.4*	323.6 ± 100.8
	HDCT	342.4 ± 91.6*	349.8 ± 90.7*	345.0 ± 90.8*	340.0 ± 91.3*	335.2 ± 92.2*	325.4 ± 93.8*
		$p = ns$					

Data are attenuation values in Hounsfield units given as mean ± SD

SDCT standard definition computed tomography, HDCT high definition computed tomography

* $p < 0.05$ for each values versus in-stent

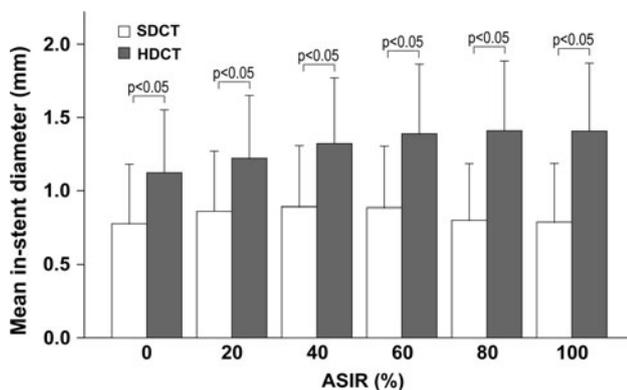


Fig. 4 The mean measured in-stent luminal diameter was significantly higher in HDCT than in SDCT ($p < 0.05$) while no significant difference in stent length was measured

0.8 ± 0.4 mm, 0 %ASIR; $p < 0.05$), which also holds true for all ASIR reconstructions. Addition of increasing ASIR contributions had no impact on the in-stent luminal diameter in SDCT (0.8 ± 0.4 mm, 100 % ASIR), while it tended to increase in HDCT (1.4 ± 0.5 mm, 100 % ASIR) although the comparison to FBP fell short of statistical significance (Fig. 4).

There was no significant difference in mean measured stent length in HDCT and SDCT (23.4 ± 8.3 mm vs. 23.8 ± 8.3 mm, 0 %ASIR; $p = ns$), which remained unaffected by ASIR. Figure 5 demonstrates a representative stent scanned in HDCT and SDCT reconstructed with all ASIR contributions.

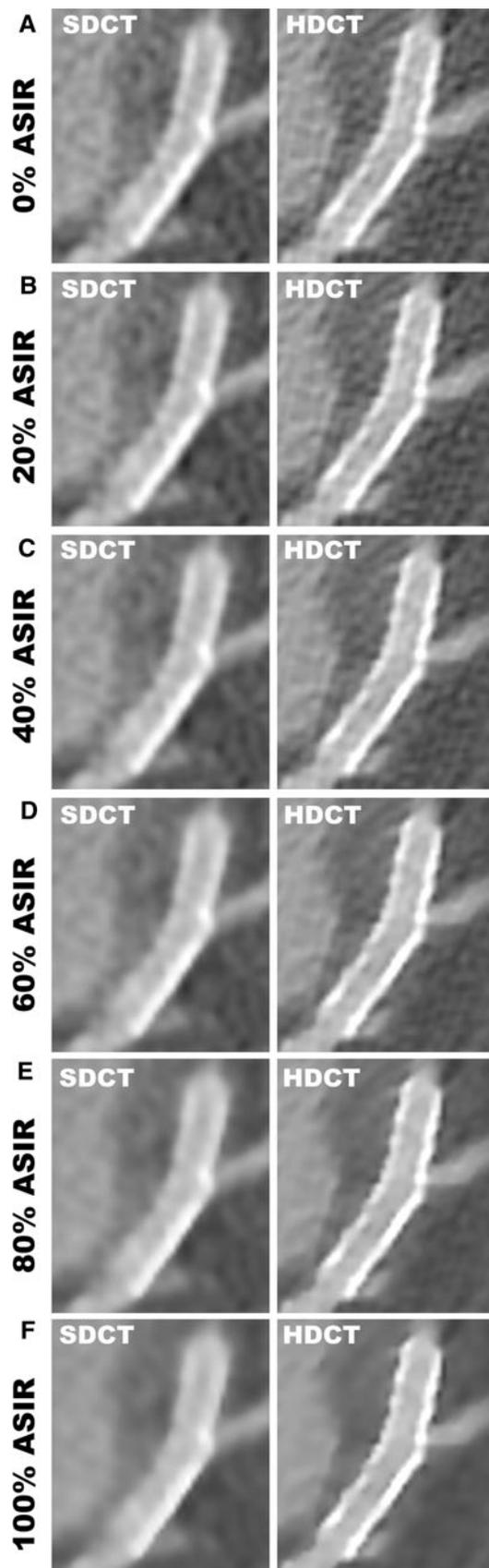
Discussion

This study reports on the first head-to-head comparison of the in vivo visualisation and quantitative assessment of coronary stents using a latest generation HDCT versus

SDCT. The main finding is the fact that partial volume effects were significantly reduced by HDCT scanning. As a result, the proportion of stents imaged successfully with no or only minor artefacts increased substantially from 44 to 96 % of stents by HDCT, underlining that partial volume effects represent a major limitation of coronary stent imaging by SDCT.

Although the rate of restenosis has substantially decreased over the past years, it remains a non-negligible drawback of coronary stenting. Therefore, a reliable non-invasive diagnostic imaging tool for differentiation of in-stent restenosis from progression of CAD in non-stented segments remains highly desirable. CCTA has rapidly evolved over the past years into a widely used alternative of conventional invasive coronary angiography supported by a growing body of literature. However, visualization and therefore robust evaluation of stents by SDCT has been hampered so far by high attenuation of the stent material interacting with the luminal contrast material attenuation due to blooming and partial volume artefacts. The latter artificially increase the measured in-stent attenuation (increase in HU) and decrease the apparent stent lumen diameter therefore affecting the image interpretation. The HDCT was designed to allow an increase in spatial resolution ($0.23 \text{ mm} \times 0.23 \text{ mm}$) by delivering more views per rotation and integrating a new detector. In our study the latter was paralleled by an increase in noise, which is in line with previous studies [16]. Therefore this technical refinement has been complemented by new reconstruction algorithms such as ASIR to compensate for the noise increase [21]. In our study in both, HDCT and SDCT scanning, image noise was significantly reduced using 40 % of ASIR or more, whereas this had no impact on the mean signal.

Although the HDCT provides a high spatial resolution in this study radiation dose was comparable to SDCT scans.



◀**Fig. 5** Curved multiplanar reconstruction of a stent scanned with 64-slice SDCT (*left column*) and HDCT (*right column*) reconstructed by 0 % ASIR (**a**), 20 % ASIR (**b**), 40 % ASIR (**c**), 60 % ASIR (**d**), 80 % ASIR (**e**) and 100 % ASIR (**f**)

As the potential carcinogenic risk of CCTA has been vividly and extensively discussed and put controversially into perspective of the potential benefits of the method [22], it is very important to note that despite using HDCT, the prospective ECG triggering protocol still allows low dose CCTA yielding a mean radiation dose (1.8 ± 0.7 mSv) comparable to or even lower than reported in other CCTA studies [12, 23–25].

In vitro stent imaging analyses have shown that artefacts from coronary stents leading to artificial luminal narrowing, due to partial volume artefact from highly attenuated stent struts vary depending on the stent material [26]. Most severe artefacts have been observed in tantalum or gold-coated stents. Therefore a large variety of different stents has been included in our study in order to represent a large variety of artefact severity.

Recent studies reported that in-stent lumen cannot be interpreted in up to 42 % of coronary stents [27, 28]. Nevertheless, recent guidelines on coronary revascularization include a recommendation for CCTA follow up in patients after unprotected left main stenting [29]. This is probably due—at least in past—to the fact that stent imaging by CCTA has improved due to advancements in CT technology, particularly after introduction of 64-slice CT. Introduction of HDCT is a further promising step to improve stent image visualization and interpretation. However, whether this may translate into higher accuracy of in-stent restenosis or stenosis severity of calcified lesions remains to be evaluated. Therefore, it may be perceived as limitation of the present study that diagnostic accuracy and in-stent luminal-diameter-measurement was not compared to invasive coronary angiography as a ground of truth reference standard. Furthermore, our analysis was not performed separately for each stent due to the limited stent number in each group. However, the fact that the results are consistent with 10 different stent types further strengthens our study.

In conclusion the use of HDCT for coronary stent imaging reduces partial volume artefacts from stents yielding improved image quality versus SDCT at a comparable radiation dose.

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Conflict of interest None declared.

Ethical Standard All procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national) and with the Helsinki Declaration of 1975 as revised in 2008 (5). Informed consent was obtained from all patients for being included in the study.

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