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Coronary artery stent geometry and in-stent contrast attenuation with 64-slice computed tomography

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e-mail: pak@usz.ch Tel.: +41-1-2553555 Fax: +41-1-2554414 **Abstract** We aimed at assessing stent geometry and in-stent contrast attenuation with 64-slice CT in patients with various coronary stents. Twentynine patients (mean age 60±11 years; 24 men) with 50 stents underwent CT within 2 weeks after stent placement. Mean in-stent luminal diameter and reference vessel diameter proximal and distal to the stent were assessed with CT, and compared to quantitative coronary angiography (QCA). Stent length was also compared to the manufacturer's values. Images were reconstructed using a medium-smooth (B30f) and sharp (B46f) kernel. All 50

stents could be visualized with CT. Mean in-stent luminal diameter was systematically underestimated with CT compared to OCA (1.60 \pm 0.39 mm versus 2.49 ± 0.45 mm; P<0.0001), resulting in a modest correlation of OCA versus CT (r=0.49; P<0.0001). Stent length as given by the manufacturer was 18.2±6.2 mm, correlating well with CT (18.5 \pm 5.7 mm; r=0.95; P < 0.0001) and OCA (17.4±5.6 mm; r=0.87; P<0.0001). Proximal and distal reference vessel diameters were similar with CT and OCA (P=0.06and P=0.03). B46f kernel images showed higher image noise (P < 0.05) and lower in-stent CT attenuation values (P < 0.001) than images reconstructed with the B30f kernel. 64-slice CT allows measurement of coronary artery in-stent density, and significantly underestimates the true in-stent diameter compared to QCA.

Keywords Stents · 64-slice CT · Coronary angiography

Introduction

Multi-detector row computed tomography (CT) is increasingly becoming an important modality for the non-invasive imaging of patients with suspected coronary artery disease (CAD). Recent studies with 16- and 64-detector row CT showed a high sensitivity and specificity for the detection of coronary artery stenoses [1–7]. On the other hand, assessment of coronary artery stents with multi-detector

row CT still remains a major problem, due to partial volume artifacts from the highly attenuated stent struts, which lead to artificial narrowing of the lumen and thus hamper reliable visualization of the stent lumen [8–14]. Stents are now used in approximately 80–90% of all percutaneous coronary interventions [15]. Although the introduction of drug-eluting stents has significantly reduced the incidence of in-stent restenosis [16], an efficient noninvasive diagnostic tool for the follow-up after stent

implantation is of great clinical interest. While accuracy of stent lumen analysis was modest with 4-, 16- and 40-detector row CT scanners, the recently developed 64-slice CT scanner may allow more accurate stent characterization, due to an increased spatial and temporal resolution [17]. First in vitro and ex vivo experience with coronary artery stents using 64-slice CT has shown a superior visualization of the stent lumen and in-stent stenosis compared with 16-detector row CT [18, 19]. The aim of our study was to prospectively assess quantitative parameters, such as stent geometry and in-stent contrast attenuation, with 64-slice CT in patients with various coronary artery stents.

Materials and methods

Patients

The study protocol was approved by the local ethics committee, and written informed consent was obtained from all patients. Between February 2005 and June 2005, we prospectively enrolled 29 patients (5 women, 24 men; mean age 60±11 years, range 39-81 years) who had undergone PCI with stent implantation. All patients and stents were examined by conventional coronary angiography with quantitative coronary angiography (QCA) within 8±14 days prior to conducting the CT scans with unchanged medication. Stent diameters and stent length measured during this invasive angiography were used for comparison with CT. Only patients in sinus rhythm able to perform breath hold commands were included. Exclusion criteria were renal insufficiency (serum creatinine level >115 µmol/L) or known allergy to iodinated contrast agents.

Multi-detector row CT protocol

All CT examinations were performed on a 64-slice CT scanner (Somatom Sensation 64, Siemens Medical Solutions, Forchheim, Germany) with a detector collimation of 32×0.6 mm, a slice collimation of 64×0.6 mm, and a table feed of 9.2 mm per rotation. Gantry rotation time was 330 ms, tube voltage was 120 kV, and effective tube current-time product 650-780 mAs. All subjects received 0.4 mg nitroglycerin sublingually 2 minutes before data acquisition. A bolus of 80 ml non-ionic, iodinated contrast agent (Visipaque 320 mg/ml, Amersham Health, Buckinghamshire, United Kingdom) was continuously injected into an antecubital vein via a 18-20-gauge catheter at a flow rate of 5.0 ml/s, followed by a 30 ml saline flush. An automated bolus tracking system was used to synchronize the arrival of the contrast material with initiation of the scan. A region of interest (ROI) was placed into the aortic root, and image acquisition started 5 s after the signal density reached the predefined threshold of 140 Hounsfield

units (HU). After this, scanning was performed covering the distance from the tracheal bifurcation to the diaphragm. For optimal heart phase selection, retrospective electrocardiography (ECG) gating was used. Axial images synchronized to the ECG were reconstructed by a single sector (<65 bpm) or two-sector (>65 bpm) algorithm, using data from one or two consecutive heart beats. The estimated effective radiation dose (using ECG-controlled dose modulation) for this protocol is in the range of 6.9–11.1 mSv [20]. Two sets of CT images were reconstructed, one with a medium-smooth kernel (B30f; 50% value of the modulation transfer, approximately 4.0 cm⁻¹), the other with a sharp kernel (B46f; 50% value of the modulation transfer, approximately 5.0 cm⁻¹). Images were reconstructed every 5% interval from 0 to 95% of the R-R interval, with an effective slice thickness of 0.75 mm at 0.5 mm increments. The images in that phase showing least motion artifacts were used for further analysis. As previously reported [21]. the optimal reconstruction phase was found at 60% and 65% of the R-R interval, with the individual %-interval presenting the beginning of the reconstruction. All data were transferred to a remote workstation (Leonardo with Syngo InSpace4D application, Siemens, Forchheim, Germany) for further post-processing.

Stent visualization and image quality

Transverse CT images and the corresponding curved multiplanar reformations (MPR) along the course of the stents were analyzed by two independent radiologists, each with 5 years of experience in cardiovascular radiology, who were both blinded to the clinical history, to the indication for referral, and to the results from invasive coronary angiography. The data sets reconstructed with the sharp kernel (B46f) were used for image quality and stent patency analysis, because they are known to provide a better delineation of the stent as compared to images reconstructed with the medium-smooth (B30f) kernel [11, 18]. The two readers were allowed to individually adjust the window settings in each patient. The stent was considered assessable when the stent lumen was visible and the contrast in the lumen could be visually evaluated. Image quality of each stent was qualitatively graded on a 4point scale as follows: 1 = no artifacts in the surrounding of the stent, 2 = minor artifacts, 3 = moderate artifacts partially obscuring the stent surrounding, and 4 = severe artifacts totally obscuring the stent lumen and vicinity leading to completely non-evaluable stents. Thus, scores 2 and 3 indicated partially non-evaluable stents. When artifacts were found (i.e., scores 2-4), the reasons for impaired image quality were assigned as partial volume artifacts, motion artifacts or calcifications. Examples for each artifact type are provided on Fig. 1. Finally, stent patency was visually graded according to the proportion of the stent lumen that showed no contrast enhancement:

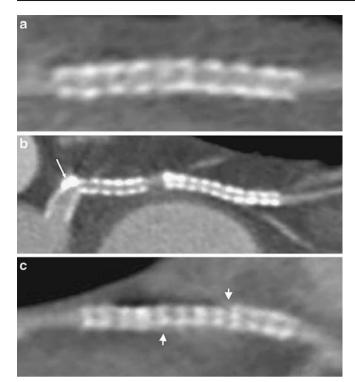


Fig. 1 Three examples of artifact types deteriorating image quality of coronary artery stents with 64-slice CT using curved multiplanar reconstruction along the stent-axis. a Partial volume artifact from the highly attenuated metal stent strut (image quality score 3), b minor artifact due to coronary vessel wall calcification (arrow) in the proximal stent (image quality score 2), and c moderate motion artifact (arrowheads) in a patient with a heart rate of 76 bpm (image quality score 3)

grade 1 = homogenous enhancement with no lumen narrowing indicating a patent stent, grade 2 = eccentric or concentric non-enhanced area between the stent and the contrast-enhanced lumen with a lumen decrease $\leq 50\%$ —indicating a non-significant in-stent restenosis, grade 3 = non-enhanced area within the stent leading to a reduction of the contrast-enhanced lumen >50%—indicating a significant in-stent restenosis, and grade 4 = no contrast enhancement within the stent indicating stent occlusion.

Attenuation measurements

Attenuation measurements were performed on transverse and curved MPR images reconstructed with both medium-smooth (B30f) and sharp (B46f) kernels by the same two radiologists in consensus. For all measurements, the images were zoomed to ensure exact placement of the ROIs. Image noise was measured by calculating the standard deviation of the mean CT attenuation in a ROI set in the ascending aorta. CT attenuation was measured within the stent lumen and in the coronary artery lumen 5 mm proximal and distal to the stent. Areas with decreased attenuation, indicating stenosis, were avoided when setting the ROI. At each site,

measurements were made on transverse images of three adjacent sections, and the results were averaged. If the axis of the stent was neither perpendicular nor parallel to the imaging plane, the measurements were performed on the curved MPR images along the course of the stent.

Stent geometry

Stent geometry was analyzed on curved MPR images reconstructed with the sharp kernel (B46f), as the B46f kernel has been shown to be superior for quantitative assessment of stent geometry with both 16- and 64-detector row CT [11, 18, 22]. For all measurements, the images were zoomed to ensure exact placement of the calipers. To standardize the analysis, images were displayed with a fixed window level at 200 HU and a window width at 1500 HU, as this window setting has been reported to provide good visualization of coronary artery stents with respect to in-stent luminal dimensions and stent strut contrast enhancement, while changes in these parameters may interfere with the results of geometry measurements [9, 11]. The in-stent luminal diameter was measured three to six times (depending on the length of the stent) by using electronic calipers, and all measurements were averaged for each stent. Similarly, stent length and diameters of the reference vessel 5 mm proximal and 5 mm distal to the stent were measured twice, and values were averaged. An example illustrating stent geometry measurements is provided in Fig. 2.

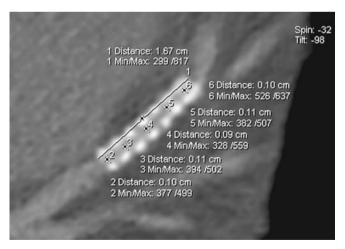


Fig. 2 Curved multiplanar reconstruction of 64-slice CT coronary angiography images illustrating stent geometry measurements. Instent luminal diameter was measured 3 to 6 times (depending on the length of the stent) by using electronic calipers and all measurements were averaged for each stent. Stent length was also measured several times and the values were averaged

Quantitative coronary angiography

Angiograms of the target vessels were obtained in at least two orthogonal projections after administering intracoronary nitrates. One cardiologist with 2 years of experience with invasive coronary angiography, who was blinded to the CT results, examined each catheter angiogram. First, the sites of stent implantation were identified before contrast injection. Thereafter, the contrast-filled angiography catheter was used as reference for calibration by using a computerized QCA analysis software (Xcelera, Philips Medical Systems, the Netherlands). Normal reference segments were identified 5 mm proximal and 5 mm distal to the stent. Mean in-stent luminal diameters, stent lengths, and reference vessel diameters were obtained on diastolicphase images in each patient. The results from two orthogonal views (average) were recorded. Narrowing of >50% of in-stent luminal diameter in relation to a reference segment was defined as significant in-stent restenosis.

Statistical analysis

Quantitative variables were expressed as mean±standard deviation. Inter-observer agreements for image quality and stent patency read-out were expressed as Cohen κ statistics and were interpreted as follows [23]: A κ value of 0.20 or less indicated poor agreement; a k value of 0.21–0.40, fair agreement; a k value of 0.41-0.60, moderate agreement; a κ value of 0.61-0.80, good agreement; and a κ value of 0.81-1.00, excellent agreement. Image noise in CT images reconstructed with B30f and B46f kernels, CT attenuation between the different measured sites, and CT attenuation between the B30f and B46f kernel CT images was compared. Pearson correlation coefficients were calculated to compare the CT attenuation in the stent lumen and in the coronary artery segments proximal and distal to the stent. Pearson correlation analysis and Bland and Altman analysis [24] were performed to compare quantitative stent geometry, and reference vessel diameter measurements between CT, QCA, and the manufacturer values. Comparisons were performed using the two-sided paired Student t-test and analysis of variance (ANOVA) using the Scheffe's F-test. A P value less than 0.05 indicated a statistically significant difference. All statistical analyses except calculation of Cohen's kappa value were performed using a commercially available software package (StatView for Windows, Version 5.0.1, SAS Institute, Cary, NC, USA). Cohen's kappa was calculated using the Statistical Package for the Social Sciences (SPSS 12.0 for Windows, SPPS Incorporation, Chicago, IL, USA).

Results

Study population

Twenty-nine consecutive patients with 50 stents (1 to 5 stents per patient, mean 1.9 ± 1.3 stents) were analyzed with 64-slice CT and QCA. Seventeen patients were studied after acute myocardial infarction. One patient had undergone prior coronary artery bypass surgery. Nineteen patients were on continuous β -blocker medication. In six patients, intravenous metroprolol (5 to 20 mg) was administered prior to the CT examination to achieve a target heart rate <75 beats per minute (bpm). The average heart rate during scan was 67 ± 12 bpm (range 40-91 bpm). The clinical characteristics of the study group are summarized in Table 1.

Stent characteristics and location

Forty-two drug eluting and eight bare metal stents were analyzed in the left anterior descending (LAD; n=23), in the left circumflex (LCX; n=8), in the right coronary (RCA; n=13) artery, and in side branches (n=6). Five different types of stents were included, i.e. Cypher Select (Cordis, Miami, FL, USA; n=28), Taxus (Boston Scientific, Boston, MA, USA; n=14), Lekton Motion (Biotronik, Switzerland; n=5), AVE (Medtronic, Natick, MA, USA; n=2), and Driver (Medtronic, Natick, MA, USA; n=1). Labeled stent di-

Table 1 Patient characteristics

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Clinical characteristics	Values			
No. of patients	29			
Age (years)	60 ± 11			
Male gender	24			
Body mass index (kg/m ²)	27±5			
Reasons for coronary angiography				
ST-segment elevation myocardial infarction	8			
Non-ST-segment elevation myocardial infarction	9			
Chest pain	12			
Coronary artery disease				
Single-vessel	12			
Two-vessel	9			
Three-vessel	8			
Previous stent angioplasty ≥6 months	10			
Previous coronary artery bypass grafting	1			
Cardiovascular risk factors				
Smoking history	13			
Hypertension	20			
Diabetes	5			

Data are numbers of patients (unless otherwise noted).

ameters were distributed as follows: 2.25 mm (n=5), 2.5 mm (n=7), 2.75 mm (n=13), 3.0 mm (n=17), and 3.5 mm (n=8). Strut thickness ranged from 0.008 to 0.14 mm. One stent (Driver) was made of a cobalt alloy, the remaining stents were made of stainless steel.

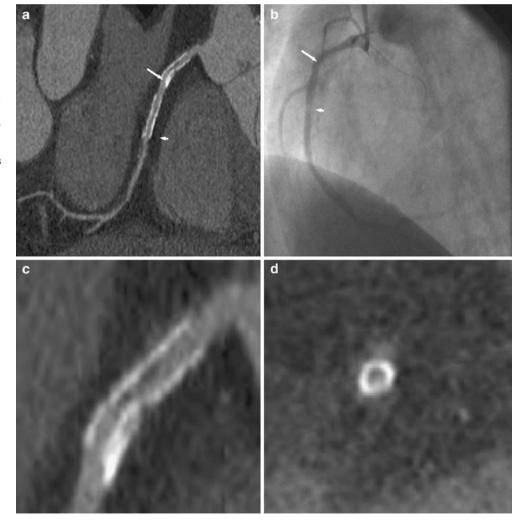
Stent visualization and image quality

All 50 stents could be visualized with 64-slice CT. Overall image quality was good, with a mean score of 1.8 ± 0.7 (inter-observer agreement κ =0.73). The score tended to be higher (2.2±0.7) for stents with a diameter <3.0 mm versus those with a diameter ≥3.0 mm (1.7±0.6; P<0.05). Using the best reconstruction interval, artifact-free images (score 1) were found in 36% of the coronary stents (n=18), minor artifacts (score 2) were found in 50% (n=25), and moderate artifacts (score 3) were present in 14% (n=7). None of the coronary artery stents was rated as being completely non-evaluable

(score 4). Reasons for impaired image quality were partial volume artifacts in 66% (n=21), motion artifacts in 25% (n=8), and calcifications in 9% (n=3). Sixty percent of the patients with motion artifacts had heart rates >70 bpm (mean 74 ± 12 bpm). Lumen interpretability did not differ significantly from one stent location to another, or from one type of stent to another.

The inter-observer agreement for stent patency grading was good (κ =0.82). Both readers correctly identified one stent in the LAD (Cypher Select) with significant in-stent restenosis (\geq 50% diameter stenosis, grade 3) and two serial stents in the RCA (2×AVE) in one patient with non-significant in-stent restenosis (grade 2) (Fig. 3). These three stents had been implanted at a previous intervention 17 and 7 months before. In two patients, the stents were incorrectly rated by one reader as having a non-significant in-stent restenosis (grade 2) and by the other reader as being patent (grade 1), and vice versa. All other stents were correctly classified by both readers as being patent (grade 1).

Fig. 3 Two AVE stents in the right coronary artery demonstrated with 64-slice CT using (a) curved multiplanar reconstruction along the vessel, and (b) by conventional coronary angiography. Proximal stent shows a non-significant in-stent restenosis (arrows) with a crescent-shaped lack of contrast enhancement (magnified parallel and perpendicular image, c and **d**). Distal stent (**a** and **b**) shows also non-significant in-stent restenosis (arrowheads), corresponding to neointimal hyperplasia



Attenuation measurements

The mean image noise measured on the B30f kernel images (22.1±6.3 HU) was significantly lower than that on the B46f kernel images (35.1 \pm 11.0 HU; P<0.05). Table 2 lists the mean CT attenuation values measured within the stent lumen and in the reference vessel lumen proximal and distal to the stent. In-stent luminal CT attenuation measured on the B46f kernel images was significantly lower compared to in-stent luminal CT attenuation measured on the B30f kernel images (P < 0.001). No significant differences in CT attenuation between the B46f kernel and B30f kernel images was found in the reference lumen proximal and distal to the stent. In-stent luminal attenuation was higher than the attenuation in the proximal and distal reference vessel (P<0.001). The in-stent luminal CT attenuation measured on the B46f kernel images was closer to the densities measured in the proximal or distal reference vessels images (F-test; all P=ns) when compared to in-stent luminal CT attenuation on the B30f kernel images (F-test: 69.8; P < 0.0001). The in-stent CT attenuation measured on both B30f kernel and B46f kernel images showed a weak but significant correlation with the attenuation in the reference vessels proximal and distal to the stent (Table 3).

Stent geometry

True average stent length as given by the manufacturer $(18.2\pm6.2 \text{ mm})$ showed an excellent correlation with 64-slice CT $(18.5\pm5.7 \text{ mm}; r=0.95; P<0.0001)$ but somewhat less with QCA $(17.4\pm5.6 \text{ mm}; r=0.87; P<0.0001)$. By contrast, mean in-stent luminal diameter was systematically underestimated with 64-slice CT compared to QCA $(1.60\pm0.39 \text{ mm} \text{ versus } 2.49\pm0.45 \text{ mm}; \text{ mean difference } -0.89 \text{ mm}; P<0.0001)$ and limits of agreement were wide (-0.03 to -1.71 mm), indicating only a modest correlation and agreement of QCA versus 64-slice CT (r=0.49;

Table 2 Comparison of contrast-enhanced CT attenuation measurements on images reconstructed with medium-smooth and sharp kernels

Location of	Measured attenuation values (HU)			
measurement	Medium smooth (B30f) kernel	Sharp (B46f) kernel	P value ^a	
Stent				
In-stent lumen	$483.7\!\pm\!143.2$	388.0 ± 96.8	< 0.001	
Reference vessel				
Proximal to stent	325.3 ± 52.6	322.1 ± 55.0	0.69	
Distal to stent	291.7 ± 73.9	296.9 ± 63.5	0.44	

Data are attenuation values as mean±SD. All absolute values given in Hounsfield units (HU).

Table 3 Correlation of contrast-enhanced CT attenuation measurements in the stent lumen and reference vessel with medium-smooth and sharp kernels

Reference vessel	In-stent lumen		
	Medium smooth (B30f) kernel	Sharp (B46f) kernel	
Proximal to stent Distal to stent	0.30 0.40	0.40 0.27	

Data are Pearson correlation coefficients (r) for bivariate analysis (P < 0.001 for all comparisons).

P < 0.0001) (Fig. 4c). Mean in-stent luminal diameter values as given by the manufacturer (2.88±0.38 mm) correlated slightly better with OCA (r=0.66; P<0.0001) than to 64-slice CT (r=0.53; P<0.0001), with somewhat narrower limits of agreement (Fig. 4a,b). Accordingly, the extent of underestimation of in-stent diameters was significantly lower between the manufacturer and QCA (mean difference, 0.38 mm) versus between the manufacturer and 64-slice CT (mean difference, 1.27 mm; P < 0.0001), whereas the F-test disclosed no significant difference for the limits of agreement between these two comparisons (F-test: 3.82; P=0.057). Quantification of vessel diameters by 64-slice CT in the reference vessel 5 mm proximal and 5 mm distal to the stent was comparable to QCA (Table 4). Quantification of stent diameter by 64-slice CT was not affected by stent size (Table 4) nor by stent type (Table 5).

Discussion

This study reports the in vivo visualization and quantitative assessment of various coronary artery stents compared to QCA. From a total of 50 examined stents, all reached sufficient image quality to allow assessment of stent lumen. Even the smallest stents could be correctly analyzed with 64-slice CT, as opposed to previous studies with 16-slice CT [25], indicating a possible improvement of the 64-slice CT technique compared to previous scanner generations. However, quantitative analysis by 64-slice CT substantially underestimates in-stent luminal diameters (by at least 35%) as compared to the gold standard QCA.

A reliable noninvasive diagnostic procedure for followup of patients with previous stent angioplasty would be desirable in symptomatic patients with suspected in-stent restenosis, as the absolute number of stent implantations increases while the restenosis rate has remained important, i.e. between 6% and 12% [16]. Although multi-detector row CT plays an increasing role as an alternative to conventional coronary angiography, reliable visualization of coronary artery stents with CT has been difficult so far, due to partial volume artifacts from the highly attenuated

^aCalculated with two-sided paired Student *t*-test.

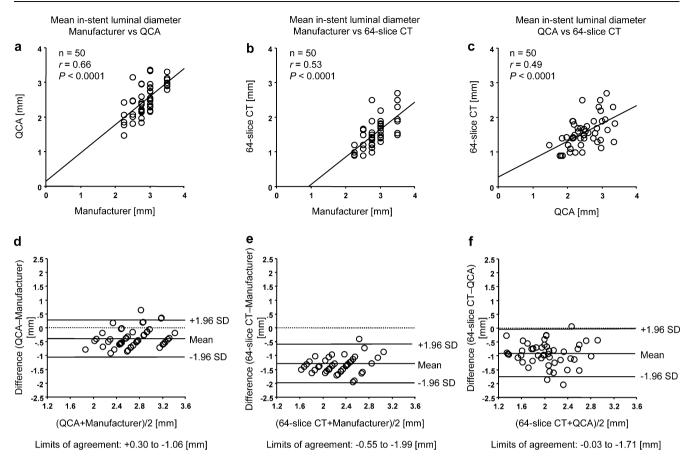


Fig. 4 Linear regression analysis for mean in-stent luminal diameter in millimeters (mm) in all stents (n=50) between (a) manufacturer and quantitative coronary angiography (QCA) (Pearson correlation; r=0.66, P<0.0001),(b) manufacturer and 64-slice CT (Pearson correlation; r=0.53, P<0.0001), and (c) between QCA and 64-slice

CT (Pearson correlation; r=0.49, P<0.0001). Bland–Altman plots for mean in-stent luminal diameter between (**d**) manufacturer and QCA, (**e**) manufacturer and 64-slice CT, and (**f**) between QCA and 64-slice CT

stent struts. In our study, none of the 50 stents was incorrectly identified with CT as having a significant stenosis. However, as only one high grade in-stent

restenosis und two stents with non-significant in-stent restenosis were found, no meaningful number on sensitivity can be provided.

Table 4 Comparison of in-stent luminal diameters and reference vessel diameters between quantitative coronary angiography (QCA) and 64-slice CT

Mean diameter	QCA	64-slice CT	Mean difference		P value ^a
			Abs.	%	
All stents $(n=50)$					
In-stent diameter	2.49 ± 0.45	1.60 ± 0.39	-0.89	-35.7	< 0.0001
Stents $\geq 3 \text{ mm } (n=25)$					
In-stent diameter	2.73 ± 0.37	1.78 ± 0.34	-0.95	-34.8	< 0.0001
Stents <3 mm $(n=25)$					
In-stent diameter	$2.24 {\pm} 0.40$	1.43 ± 0.36	-0.81	-36.2	< 0.0001
Reference vessel					
Proximal to stent	2.24 ± 0.61	2.49 ± 0.61	+0.25	+11.2	0.06
Distal to stent	$1.86 {\pm} 0.59$	2.15 ± 0.63	+0.29	+15.6	0.03

Data shown as mean±SD. All absolute values given in mm. ^aCalculated with two-sided paired Student t-test.

Table 5 Comparison of in-stent luminal diameters between quantitative coronary angiography (QCA) and 64-slice CT according to stent type

Mean stent diameter	QCA	64-slice CT	Mean difference		P value ^a
			Abs.	%	
Cypher $(n=28)$					
In-stent diameter	2.40 ± 0.45	1.61 ± 0.23	-0.79	-32.9	< 0.0001
Taxus $(n=14)$					
In-stent diameter	$2.55 {\pm} 0.40$	1.60 ± 0.49	-0.95	-37.3	< 0.0001
Lekton Motion $(n=5)$					
In-stent diameter	3.00 ± 0.19	1.50 ± 0.23	-1.50	-50.0	< 0.005
AVE $(n=2)$					
In-stent diameter	2.10 ± 0.39	1.21 ± 0.30	-0.89	-42.4	NA^b
Driver $(n=1)$					
In-stent diameter	3.36	1.83	-1.53	-45.5	NA^b

Data shown as mean±SD. All absolute values given in mm. ^aCalculated with two-sided paired Student *t*-test. ^bNA=not applicable.

In vitro analyses with CT of various stent types with different diameters have shown that artifacts depend on strut thickness, which leads to artificial narrowing and thus reduced interpretability of the intra-stent lumen. Furthermore, artifacts vary substantially according to the metal employed, the problem being most severe with tantalum or gold-coated stents [22, 26, 27]. In a study published by Krueger et al. [8] a total of 32 stents were assessed in vivo by 4-detector row CT. The lumen of the stents could not be interpreted in any of these stents due to severe artifacts. This finding was independent of the size and type of examined stents. Gilard et al. [28] analyzed 232 stents in vivo by 16-detector row CT. Intra-stent lumen was interpretable only in about one half of the patients. A stent diameter of 3 mm or more was associated with better lumen visibility and better ability to detect intra-stent restenosis. In a recent study using 40-detector row CT, Gaspar et al. [29] assessed 111 stents in 65 patients referred for repeat invasive angiography after clinical suspicion of in-stent restenosis. Although in-stent restenosis could be diagnosed with moderate sensitivity (88.9%) and specificity (80.6%), the high negative predictive value (97.4%) implies a significant role for multi-detector row CT in excluding in-stent restenosis.

The 64-slice CT scanners provide a high temporal and spatial resolution, potentially reducing partial volume artifacts in stent imaging [17]. Therefore, the introduction of 64-slice CT might improve the yield in this application. Our results confirm the progress achieved by the latest technique, as in the present study each stent lumen was at least partially assessable. Mean in-stent luminal diameters assessed by 64-slice CT were smaller than those measured on conventional angiograms. These findings are in accordance with a previous study by Hong et al. [11] where a roughly 20% size underestimation of the mean instent luminal diameters was also reported. This indicates

that the present spatial resolution of 64-slice CT may still be not sufficient to avoid partial volume averaging artifacts. This holds true, although the role of QCA as gold standard is challenged by the fact that true stent length as given by the manufacturer showed a better agreement with 64-slice CT than with QCA. The underestimation of in-stent luminal diameters reflecting the problem of partial volume averaging close to the stent strut may be also the reason for the two incorrect ratings of non-significant in-stent restenoses in two patients with patent stents.

Similar to the 16-detector row CT study by Hong et al. [11], we have demonstrated the capability of 64-slice CT to assess stent patency by means of intraluminal contrast enhancement measurements. Because of partial volume averaging artifacts from the stent struts, the CT attenuation values were significantly higher than those measured in the reference coronary vessel proximal and distal to the stent. The in-stent luminal CT attenuation measured on images reconstructed with the sharp kernel was closer to that measured in the proximal or distal lumen than was the instent luminal CT attenuation measured on the medium-smooth kernel reconstructed images. This can be explained by a reduction in partial volume averaging artifacts with the sharp convolution kernel reconstructions [11].

Our study has several limitations. As the study was primarily designed to assess stent geometry, 64-slice CT was performed within a few days after the actual stent implantation. Only 15 stents had been implanted at a previous intervention, and therefore just a small number of stents with restenosis was found. As a consequence, no meaningful data on sensitivity and specificity for the detection of in-stent restenosis by 64-slice CT can be provided. Another important limitation is that the true gold standard for assessment of stent geometry would have been intravascular ultrasound (IVUS) which has been shown to characterize stent geometry more accurately than invasive

angiography [30]. In our study, mean in-stent luminal diameter values as given by the manufacturer were also slightly underestimated by QCA. A possible reason for this might be related to incomplete stent expansion. Nakamura et al. [31] demonstrated that, despite excellent angiographic appearance, the majority of stents deployed at low pressure remain suboptimally expanded by IVUS criteria. Potential explanations for this persistent stent undersizing measured by QCA include balloon underexpansion (despite high inflation pressures), acute stent recoil, or a combination of the two mechanisms. Finally, effective radiation doses were not individually calculated in each patient, but derived from estimates from commercially available software.

In conclusion, 64-slice CT allows measurement of coronary artery stent density, and significantly underestimates the true in-stent diameter compared to QCA.

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