Quantitative analysis of epicardial fat volume: effects of scanning protocol and reproducibility of measurements in non-contrast cardiac CT vs. coronary CT angiography

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Background: Several studies have focused on the role of epicardial fat in the pathogenesis of cardiovascular disease (CVD). The main purpose of the study was to evaluate a computerized method for the quantitative analysis of epicardial fat volume (EFV) by non-contrast cardiac CT (NCT) for coronary calcium scan and coronary CT angiography (coronary CTA).

Methods: Thirty patients (61±12.5 years, 73% male, body mass index (BMI) =25.9±6.3 kg/m²) referred to our Institution for suspected coronary artery disease (CAD) underwent NCT and coronary CTA. Epicardial boundaries were traced by 2 experienced operators (operator 1, operators 2) on 3 and 6 short-axis (SA) slices. EFV was computed with a semi-automatic method using an in-house developed software based on spherical harmonic representation of the epicardial surface. In order to analyze the inter-observer variability both the Coefficient of Repeatability (CR) and Intra Class Correlation (ICC) were computed.

Results: The total EFV was 103.62±50.97 and 94.96±67.91 cc in NCT and coronary CTA with non-significant difference (P=0.292). CR error was 10.22 cc for operator 1 and 11.31 cc for operator 2 in NCT and 7.99 cc for operator 1 and 7.75 cc for operator 2 in coronary CTA. To analyze the inter-observer variability CR and ICC were computed. CR was 8.17 and 8.39 cc with NCT and 7.07 and 7.21 cc with CTA for 6 and 3 SA slices respectively. ICC values >0.99 were obtained in all cases. The right ventricular EFV was 67.23±31.4 and 57.41±34.3 cc for NCT and coronary CTA respectively; the corresponding values for left ventricular EFV were 38.01±19.1 and 35.27±25.9 cc.

Conclusions: Both NCT and coronary CT can be used with low intra- and inter-observer variability for computer-assisted measurements of EFV. Cardiac CT may allow a fast and reliable computation of EFV in clinical setting.

Keywords: Cardiovascular disease (CVD); coronary calcium scan; coronary CT angiography (coronary CTA); epicardial fat volume (EFV); pericardial fat volume

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**Introduction**

Several studies have focused on the role of the mediastinal (intra- and extra-pericardial) fat in the pathogenesis of cardiovascular disease (CVD), due to the direct contact with the coronary vessels (1,2). Indeed, in case of excess of cardiac fat deposits the delivery of free fatty acids increases expression of the molecules promoting inflammation, which may, in turn, accelerate the atherosclerosis process (3). Although measurements of abdominal visceral fat would be the real effective CVD risk predictor, a standardized quantitative method of measurement is not used in clinical practice. In a recent study it was reported that epicardial fat volume (EFV) was the best predictor for angiographic disease severity compared to waist circumference or visceral abdominal fat volume (4). Furthermore, it has been demonstrated that vascular aging and subclinical atherosclerosis, as revealed by carotid stiffness and intima-media thickness, are related to epicardial fat thickness better than waist circumference in hypertensive patients (5). Even in non-obese patients epicardial fat may be more related to the development of coronary atherosclerosis than the traditional cardiovascular risk factors and abdominal visceral adipose tissue (6). Several imaging techniques such as echocardiography, computed tomography (CT) and magnetic resonance (MR) (7-9) have been used to measure intra- and extra-pericardial fat. The high spatial resolution of the CT may allow an accurate measurement and quantification of EFV, but specific applications have not been developed until now.

The purposes of the study were to evaluate the performance and inter- and intra-observer variability of a computerized method for the quantitative analysis of EFV and epicardial fat density (EFD) from images obtained by coronary calcium scan or non-contrast cardiac CT (NCT) and coronary CT angiography (coronary CTA).

**Methods**

**Patient population**

Our study was a retrospective analysis and included a set of 30 patients (mean age =61±12.5 years, 73% male, mean body mass index (BMI) =25.9±6.3 kg/m²) referred to our Institution for suspected CAD between April 2010 and February 2011 (Table 1), who underwent NCT scan for calcium score evaluation followed by coronary CTA for assessment of coronary arteries patency. Clinical and biochemical analysis were collected. All patients gave written informed consent for retrospective use of their data.

**Cardiac CT**

The images were acquired using a multidetector row CT (MDCT, LightSpeed VCT 64, GE Healthcare, Milwaukee, WI, USA). The following parameters were used for coronary calcium scan: detector collimation =2.5 mm, reconstruction interval =2.5 mm, detector coverage =20 mm, (8 detectors × 2.5 mm), gantry rotation time =0.35 ms, tube current =300 mA (automated modulated), and voltage =100 kV, retrospective ECG gating.

For coronary CTA the following parameters and techniques were used: retrospective ECG gating with spiral technique, 64 channel detectors along the z-axis, scan FOV 15–21 cm (depend on the patient size) from the ascending aorta to the level of the diaphragm, matrix 512x512, detector collimation 0.625 mm; reconstruction 0.6x0.6 mm, range of helical pitch 0.18–0.24, gantry rotation time 0.35 ms, tube current range =250–600 mA, depending on patient size and with automated modulation of the kV based on the BMI, 100 or 120 kV. Nonionic iodinated contrast medium (Iomeprol 400, Bracco Imaging Italia s.r.l.) was injected via a peripheral vein with a triphasic protocol using a programmable injector (Nemoto Dual Shot Injector, Nemoto Kyorindo Co. Ltd., Japan) with a two-way syringe system. In the first phase, a rapid injection of contrast medium (8 mL/sec) was performed. In the second phase, 10 mL of contrast at 1 mL/sec were injected.

<table>
<thead>
<tr>
<th>Characters</th>
<th>Values</th>
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<td>6</td>
</tr>
<tr>
<td>Age (years)</td>
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</tr>
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</tr>
<tr>
<td>Diabetes</td>
<td>7</td>
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<tr>
<td>Smokers</td>
<td>9</td>
</tr>
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<tr>
<td>Hypercholesterolemia</td>
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<tr>
<td>BMI (kg/m²)</td>
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<tr>
<td>Angina</td>
<td>19</td>
</tr>
<tr>
<td>Other symptoms (dyspnea)</td>
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Table 1 Patients characteristics
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simultaneously with 25 mL of saline at 2.5 mL/sec; and in the third phase 35 mL saline flush was administered at 4 mL/sec (maximal total volume of contrast medium 110 mL). At the time of the scan, a region of interest was placed in the right ventricular cavity to detect the pre-assigned peak of contrast enhancement, typically 150 Hounsfield Unit (HU). Scans were performed during breath-hold; patients were monitored continuously through single-lead electrocardiography. As clinically indicated, patients were pre-treated with a β-blocker (metoprolol) up to 5 mg intravenously to lower the heart rate to a value below 65 beats per minute (bpm), and with a nitrate (isosorbide dinitrate) up to 1 mg intravenously to determine vasodilatation just prior to CT imaging.

**EFV measurement**

The images acquired for NCT and coronary CTA were reconstructed with a slice thickness of 2.5 mm and slice distance of 2.5 mm without overlap. DICOM images were transferred to a research workstation for fat quantification. Users were asked to trace the pericardial contours (15 points at least per boundary) in multi-planar reformat (MPR) planes, corresponding to 2- and 4-chamber views, and short axis (SA) views placed at the ventricular base, at the level of the papillary muscles (mid-ventricular region) and at the level of the apex. Boundaries of the computed surface were shown superimposed on the CT images (Figure 1). During the operation, a spline-interpolated curve was shown allowing the user to evaluate the adherence of provided data with the anatomical boundary. In this way, tracing complete curves was avoided which resulted in time saving without significant loss of accuracy. It was possible to draw additional contours in intermediate slices. Furthermore, the

![Figure 1](image-url) Epicardial fat volume (EFV) measurement: segmentation method. The boundaries of the pericardium were traced in 2 multiplanar reformat (MPR) planes, corresponding to 4- and 2-chamber views, and in 3 or 6 slices in short axis view (red lines with green dots). Segmented boundaries of pericardium were shown to the user as superimposed yellow line on short axis slices. Then the user was asked to split the cardiac region into left and right zones by tracing an interventricular plane in two short axis slices.
procedure allowed the user to split the cardiac region into left and right zones by tracing an inter-ventricular plane in 2 SA views. The superior portion of the pericardial space above the atrioventricular sulcus that included atria and mediastinal vessels was excluded because the computation was very complex and would have required strong manual intervention.

Epicardial surface was computed by using a method previously developed at our Institution (10) that relies on the properties of spherical harmonics (11), which are able to represent any continuous surface of the sphere. Smooth surfaces, such as the epicardial one, can be represented by a reduced number of the terms in the series, resulting in a light computational load. The combination coefficients were estimated by minimizing the distance between the points of the traced contours and the surface itself.

Local manual editing/refinement and re-drawing the boundaries were possible however it was required only in a few cases. EFV was computed from the segmented regions by counting the voxels with a gray level in the range of (−30, −190) HU. This has been validated by previous investigators (12). Results were provided in cubic centimeters (cc). The average gray level and the related standard deviation were also computed.

Experiments were focused to assess the effect of different numbers of SA slices along with the observer variability. Two trained technologists (O1 and O2) having >5 years of experience in cardiac CT and coronary CTA procedures analyzed the images of all patients twice with a time interval between the measurements of at least 2 weeks and with a random order in the presentation of images, with a total of 120 examinations for each operator. The observers were pre-trained to the use of program. In addition, in order to ensure a uniform consensus-based operation, during the training phase, the observers were asked to analyze the traced boundary and the resulting surface.

Statistical analysis
Statistical analyses were performed using Statplus® software. Observer variability was analyzed by Bland-Altman (BA) method (13). Intra-class correlation (ICC) was also evaluated. Statistical correlations were analyzed by Pearson correlation coefficient.

All continuous variables were expressed as median ± inter-quartile range. Differences between samples were tested by the Wilcoxon test. A P value <0.05 was considered statistically significant.

Results
To test the performance of the EFV software, we used CT images from a set of 30 patients referred to our institution for suspected CAD. For each scan, post-processing time took around 2–3 minutes and average computation time was about 10 seconds. The total EFV was 103.62±50.97 and 94.96±67.91 cc in NCT and coronary CTA images respectively, with non-significant difference (P=0.292). Non-significant difference was also observed between EFV computed using 3 and 6 SA slices. A close correlation between EFV measured in NCT and coronary CTA images was found in both operators. EFV measured in coronary CTA was slightly lower than in NCT, the behavior was virtually the same using 3 and 6 SA slices (Figure 2).

According to Bland and Altman (13), we quantified the intra-observer variability through the Coefficient of Repeatability (CR), which is defined as 1.96 times the standard deviation of the differences between 2 reiterations of measurement. Therefore, difference between 2 observations lies within the range of (−CR, CR) with a probability of 95%. The lower values indicated better repeatability.

When using 3 SA slices, CR error was 10.22 cc for operator-1 and 11.31 cc for operator-2 in NCT, and 7.99 cc for operator-1 and 7.75 cc for operator-2 in coronary CTA (Figure 3). In case of 6 SA slices, CR was 6.33 cc for operator-1 and 6.93 cc for operator-2 in NCT, whereas was 6.07 cc for operator-1 and 6.14 cc for operator-2 in coronary CTA scan (Figure 4).

To analyze the inter-observer variability CR and ICC were computed. CR was 8.17 and 8.39 cc in NCT and 7.07 and 7.21 cc in CTA for 6 and 3 SA slices respectively (Table 2). ICC values >0.99 were obtained in all cases (Table 3).

The right ventricular EFV was 67.23±31.4 and 57.41±34.3 cc, in NCT and coronary CTA respectively; the corresponding values for left ventricular EFV were 38.01±19.1 and 35.27±25.9 cc. The average fat density was −86.6±6.8 HU in NCT vs. −85.4±6.2 HU in CTA, (P<0.04). The range of fat density was (−99.56, −74.1) HU in NCT vs. (−94, −75.82) HU in coronary CTA.

Furthermore, EFV showed a very close correlation with BMI (rho =0.783, P=0.0001), more influenced by the body weight than by the height (Table 4).

Discussion
A positive correlation between EFV and coronary plaque
Figure 2 Regression analysis shows excellent correlations of epicardial fat volume (EFV) measured in cubic centimeters (cc) in non-contrasted cardiac CT (NCT) and coronary CT angiography (CTA) scans in operator O1 and O2, using 3 and 6 short axis slices. Non-significant difference between EFV measured in NCT and coronary CTA was found.

burden has been reported in the literature. Also the negative consequences of increased fat deposits in the extra- or intra-pericardial regions and inside the heart (intra- and inter-myocyte cells) on the coronary circulation and on the cardiac function have been reported by previous researchers (14-16). Epicardial and mediastinal adipose tissue can mechanically and functionally compromise the systolic and diastolic function of the cardiac ventricles (17). These evidences explain the importance of a quantitative measurement of the fat volume and the efforts in obtaining tools for diagnostic interpretation and monitoring the relationship between visceral fat and atherosclerotic process.

In this paper, we used a method for semi-automatic EFV measurements, aiming to improve the accuracy of the non-standardized methods generally reported in literature. The latter were limited to manually outlined regions of interest (ROI) and preset density thresholds range for fat tissue, which was subject to inter- and intra-observer and inter-scan variability (6).

The total EFV was 64.38±46.04 and 103.62±50.97 cc (P<10^-6) in coronary CTA and NCT, respectively. Those results are similar to those published in the literature, which range from 68±34 to 124±50 mL in population-based studies (18,19) and 110±41 mL in women and 137±53 mL in men in a study including patients from the Framingham cohorts (20).

The average computation time of EFV with our software was about 10-15 seconds after tracing pericardial boundaries, and the overall time needed to complete the measurement was about 2 minutes, while others
described procedures that required more time and were thus considered to be excessively time and labor intensive (15,21,22).

To our knowledge, a comparison between measurements of EFV and epicardial fat density (EFD) in cardiac CT with and without CM was not done previously. The results of our study showed a low intra- and inter-observer variability of computer-assisted measurements of EFV in both NCT and coronary CTA, which was similarly found in previous studies (ICC of 0.95) (23,24).

Moreover intra- and inter-observer variability of the computer-assisted measurement of EFV slightly improved with CM. This may be explained by the fact that the contrasted cavities and enhanced vessels allowed a better visualization of the pericardial boundaries (Figure 5). Moreover, in NCT, there may be difficulties to identify the pericardium in lean individuals with less pericardial fat (19).

In addition, by increasing the number of SA slices from 3 to 6, it was possible to slightly decrease observer variability.

The average fat density was slightly lower in the series without CM than with CM. Furthermore, fat density range was larger in the series without CM than with CM. Partial volume effect may account for the slight observed differences between measurement done in scan with and without CM. It occurs whenever a structure is only partly within the imaging section (voxel), thereby the densities of

Figure 3 Using 3 short-axis slices, the intra-observer repeatability coefficient (CR) error in non-contrasted cardiac CT (NCT) was 10.22 cc in operator O1 and 11.31 cc in operator O2, whereas it was 7.99 cc in O1 and 7.75 cc in O2 in coronary CT angiography (CTA).
Using 6 short-axis slices, the intra-observer repeatability coefficient (CR) error in non-contrasted cardiac CT (NCT) was 6.33 cc in operator O1 and 6.93 cc in operator O2, whereas it was 6.07 cc in operator O1 and 6.14 cc in operator O2 in coronary CTA (CTA).

<table>
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<tr>
<th>Method</th>
<th>CR</th>
<th>95% CI</th>
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<tr>
<td>NCT 6 SA slices</td>
<td>8.17</td>
<td>(7.627, 8.632)</td>
</tr>
<tr>
<td>NCT 3 SA slices</td>
<td>8.39</td>
<td>(7.833, 8.865)</td>
</tr>
<tr>
<td>CTA 6 SA slices</td>
<td>7.07</td>
<td>(6.60, 7.47)</td>
</tr>
<tr>
<td>CTA 3 SA slices</td>
<td>7.21</td>
<td>(6.731, 7.618)</td>
</tr>
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SA, short-axis.

<table>
<thead>
<tr>
<th>Method</th>
<th>ICC</th>
<th>95% CI</th>
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<tr>
<td>NCT 6 SA slices</td>
<td>0.9980</td>
<td>(0.9958, 0.9990)</td>
</tr>
<tr>
<td>NCT 3 SA slices</td>
<td>0.9979</td>
<td>(0.9956, 0.9990)</td>
</tr>
<tr>
<td>CTA 6 SA slices</td>
<td>0.9995</td>
<td>(0.9992, 0.9998)</td>
</tr>
<tr>
<td>CTA 3 SA slices</td>
<td>0.9994</td>
<td>(0.9978, 0.9995)</td>
</tr>
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SA, short-axis.
Table 4 Statistical analysis univariate correlation

<table>
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<tr>
<th>Variables</th>
<th>Pearson rho</th>
<th>P</th>
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<tr>
<td>Left EFV versus BMI</td>
<td>0.787</td>
<td>8.38×10⁻⁶</td>
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<tr>
<td>Right EFV versus BMI</td>
<td>0.60133</td>
<td>2.41×10⁻⁶</td>
</tr>
<tr>
<td>EFV versus BMI</td>
<td>0.785</td>
<td>2.81×10⁻⁷</td>
</tr>
<tr>
<td>EFV versus weight</td>
<td>0.7503</td>
<td>1.81×10⁻⁶</td>
</tr>
<tr>
<td>EFV versus height</td>
<td>0.3675</td>
<td>0.0457</td>
</tr>
</tbody>
</table>

The significant values of “P” are in bold. A close correlation was found between epicardial fat volume (EFV) and body mass index (BMI), maintained both in the right EFV and left EFV, a relation more influenced by weight than by height.

Figure 5 The presence of the contrasted cavities and especially of enhanced vessels (left image) might allow a better visualization of the pericardial boundaries with respect to non-contrast CT scan (right image).

the structure and the adjacent or surrounding structures are averaged.

Furthermore, the software for epicardial fat analysis permits a systematic evaluation of total epicardial fat burden, and the segmentation between right and left ventricle. As expected, the EFV values corresponding to the right ventricle were larger than the left one, which is as expected according to the anatomical features.

The analysis of the data showed a close correlation between EFV and BMI, a relationship more influenced by weight than by height, linking EFV with other visceral fat depots. Moreover, the association between EFV and BMI was maintained in both the right EFV and left EFV, which indicate a non-selective accumulation of epicardial fat between the ventricles with the increase in body weight.

The limitation of the study is that population was small (30 patients), with a majority of males (73%) generally overweight (BMI 25.9±6.3).

Conclusions

CT may allow a fast and reliable quantification of EFV in clinical setting, a potential additional predictor of prognosis in patients with CVD. Both NCT and coronary CTA can be used to measure EFV with low intra- and inter-observer variability.

Acknowledgements

None.

Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

Disclaimer: The method for determining the volume of epicardial fat from CT images used in this study was

**Ethical Statement:** The protocol for the research project has been approved by a suitably constituted Ethics Committee of the institution within which the work was undertaken and that it conforms to the provisions of the Declaration of Helsinki (as revised in Edinburgh 2000). The subject involved in the study gave informed consent.

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