A new measurement site for echocardiographic epicardial adipose tissue thickness and its value in predicting metabolic syndrome

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Abstract

Background. Echocardiographic epicardial adipose tissue (EAT) thickness is defined as the thickness of the low-echoic area on the free wall of the right ventricle in the parasternal long-axis and short-axis views. Recent studies have suggested that it might support current risk stratification strategies in identifying an increased risk of metabolic syndrome.

Objectives. The aim of this study is to explore a new measurement site which can better reflect EAT thickness and to assess its value in predicting metabolic syndrome.

Material and methods. A total of 975 Chinese adults were measured for EAT thickness on the right ventricular anterior free wall (EAT-rv) and on the anterior interventricular groove (EAT-ivg) with echocardiography. The correlation between EAT thickness and metabolic syndrome was analyzed, as was the agreement between epicardial adipose volume (EAV) and EAT thickness. Independent risk factors of EAT thickness were identified and the predictive value of EAT thickness was assessed.

Results. Epicardial adipose tissue thickness was higher in older participants and those with obesity, diabetes, hypertension, hypertriglyceridemia, and metabolic syndrome, and it was lower in male participants. The EAT-ivg was higher in the participants with hypo-high-density-lipoprotein cholesterolemia than in those without the disorder, but the EAT-rv values were not statistically different. The kappa value was 0.524 between EAT-rv and EAV, and 0.783 between EAT-ivg and EAV. Advanced age, large waist circumference and female gender were independent risk factors of high EAT-ivg, while high-density-lipoprotein (HDL) cholesterol was a protective factor. The EAT-ivg was associated with metabolic syndrome. The area under the curve of EAT-ivg applied in predicting metabolic syndrome was greater than that of EAT-rv (0.715 vs 0.648).

Conclusions. The EAT-ivg was more consistent with EAV than EAT-rv, was independently associated with metabolic syndrome and had a higher value in predicting metabolic syndrome than EAT-rv. Therefore, the anterior interventricular groove can serve as a new measurement site which better reflects EAT thickness.

Key words: metabolic syndrome, echocardiography, epicardial adipose tissue, consistency, predictive value
Introduction

Epicardial adipose tissue (EAT) is the visceral thoracic fat depot that surrounds the heart, located between the myocardium and the visceral pericardium. It has been reported that EAT is associated with multiple pathological states, including metabolic syndrome, T2DM, atrial fibrillation, and coronary artery disease. Recent studies have shown that EAT thickness measured with echocardiography may play a limited additional role supporting current risk stratification strategies in identifying individuals at an increased risk of metabolic syndrome. Moreover, measurements of EAT thickness attained by echocardiography and computed tomography (CT) are inconsistent. This fact may be explained by the poor distribution of EAT at the measurement site. Echocardiographic EAT thickness has been defined as the thickness of the low-isoechoic area on the free wall of the right ventricle in the parasternal long-axis and short-axis views, but the distribution of EAT is unbalanced and is concentrated primarily in the interventricular and atrioventricular grooves rather than in the free wall of the right ventricle. Therefore, there is clinical value in finding a new measurement site for echocardiographic EAT thickness which can better reflect the risk of metabolic syndrome. In addition, Salami et al. have confirmed that there are significant racial differences in the distribution of EAT.

A large-sample investigation of EAT thickness in Chinese adults has not yet been published. In this study, a total of 975 Chinese adults were measured for EAT thickness on the right ventricular anterior free wall (EAT-rv) and on the anterior interventricular groove (EAT-ivg). Distribution of EAT and its consistency with epicardial adipose volume (EAV) were described, and its independent association with metabolic syndrome and its value in predicting metabolic syndrome were analyzed. The aims were to explore a new measurement site which can better reflect EAT thickness and to assess its value in predicting metabolic syndrome.

Material and methods

Participants

A total of 986 Chinese adults receiving a general check-up at the Health Management Center of Shandong Provincial Hospital between January 2017 and September 2017 were enrolled in the study. The exclusion criteria consisted of the following: 1) age <18 years or >75 years; 2) atrial fibrillation; 3) serum creatinine level >2 mg/dL; 4) hypothyroidism; 5) current or past use of glaucon-like peptide (GLP)-1, SGLT2 inhibitors, thiazolidinediones, fibrates, statins, or insulin; 6) coronary heart disease; and 7) advanced malignant tumors. Transthoracic 2-dimensional echocardiography was performed in all participants, and 11 participants were excluded because of inadequate image quality. This study was approved by the ethics committee of Shandong Provincial Hospital, China (approval No. 2016024138), and all participants provided their written informed consent.

Measurement of EAT thickness

Transthoracic 2-dimensional echocardiography was performed with a Philips iU22 ultrasound system and an s5-1 matrix-array transducer (Philips Healthcare, Bothell, USA) according to the recommendations of the European and American Societies of Echocardiography. All participants were placed in the left lateral decubitus position. Ultrasonic images were stored in specialized workstations where EAT thickness was measured by the same reader. According to recommendations by Iacobellis et al., EAT thickness on the anterior ventricular free wall (EAT-rv) was measured as shown in Fig. 1. In addition, EAT thickness on the anterior interventricular groove (EAT-ivg) at the midchordal level and the tip of the papillary muscle from the parasternal short-axis view was measured as a new measurement site, as shown in Fig. 2. The average value of 3 cardiac cycles for each echocardiographic view was used for analysis.

Quantification of epicardial adipose volume

Among these 975 participants, 120 were randomly selected for cardiac dual-source computer tomography (DSCT). Cardiac DSCT was performed using a 128-slice DSCT scanner (Siemens AG, Munich, Germany). Image reconstructions were performed at 75% RR intervals. A density ranging from −30 to −190 Hounsfield units was employed to identify adipose tissue. The pericardium was manually traced from the right pulmonary artery to the diaphragm in order to determine a region of interest. Epicardial adipose volume was assessed using a dedicated workstation (Syngo.via v. VB10A; Siemens) by the same experienced CT diagnostic physician.

Diagnosis of metabolic syndrome

Metabolic syndrome was determined according to the recommendations of the Chinese Diabetes Society, including: 1) obesity, i.e., body mass index (BMI) ≥25 kg/m²; 2) fasting blood glucose level ≥6.1 mmol/L or the use of antidiabetic medication; 3) triglyceride level ≥1.7 mmol/L; 4) high-density-lipoprotein (HDL) cholesterol count <1.04 mmol/L; and 5) blood pressure higher than 130/85 mm Hg or the use of hypotension medication. Metabolic syndrome was diagnosed when a patient had at least 3 of the above 5 components.

Statistical analysis

All analyses were performed using SPSS software v. 22.0 (IBM Corp., Armonk, USA). Continuous variables with
normal distribution were expressed as means ± standard deviations (SDs) and those without normal distribution as 25th–75th percentiles (P25–P75). The EAT thicknesses were compared between groups using the Kruskal–Wallis test. Non-conditional logistic regression analysis was performed to determine the association between metabolic syndrome and EAT thickness. The EAV and EAT thickness were grouped according to tertiles, and consistency was determined with κ statistics. Ordered logistic regression was used to identify independent risk factors of EAT thickness (quartiles). The predictive value of EAT thickness was evaluated with a receiver operating characteristic (ROC) curve, and the areas under the curve (AUC) were compared with the Z-test. Significance was set at p < 0.05.
Results

Distribution of EAT thickness

The mean age of all participants (n = 975) was 55.58 ±11.78 years; their EAT-rv was 3.01 (2.09–3.90) and their EAT-ivg was 5.27 (4.01–6.86). EAT-ivg was greater than EAT-rv (p < 0.05). The distribution of EAT thickness in all participants is shown in Table 1. Epicardial adipose tissue thickness was higher in older participants and in those with obesity, diabetes, hypertension, hypertriglyceridemia, and metabolic syndrome; it was lower among the male participants. However, EAT-ivg was higher in participants with hypo-high-density-lipoprotein cholesterolemia than in those without this condition, and EAT-rv was not statistically different between the participants with and without hypo-high-density-lipoprotein cholesterolemia.

Consistency between EAT thickness and EAV

The κ value between EAT-rv and EAV was 0.524 (p < 0.05), and that between EAT-ivg and EAV it was 0.783 (p < 0.05) (Table 2 and 3).

Risk factors of EAT-ivg

As shown in Table 4, ordered logistic regression analysis showed that advanced age, large waist circumference and female gender were independent risk factors of high EAT-ivg, and HDL cholesterol was a protective factor. On the other hand, BMI, triglyceride, HDL cholesterol, uric acid, creatinine, estimated glomerular filtration rate (eGFR), and Hba1c levels, hypertension, diabetes, and lifestyle (including eating habits, smoking and exercise habit) were not statistically significant.

Association between EAT thickness and metabolic syndrome

As shown in Table 5, non-conditional logistic regression analysis revealed that EAT-ivg was associated with metabolic syndrome, and the crude, age-adjusted,
Age- and waist-circumference-adjusted and age-, waist-circumference- and BMI-adjusted odds ratios (ORs) were 1.628, 1.529, 1.363, and 1.331, respectively. The EAT-rv was associated with metabolic syndrome when only age was adjusted for.

**Value of EAT thickness applied in predicting metabolic syndrome**

As shown in Fig. 3, the AUC of EAT-rv applied in predicting metabolic syndrome was 0.648 (standard error: 0.018), and the AUC of EAT-ivg was 0.715 (standard error: 0.017). A Z-test showed that the AUC of EAT-ivg was greater than that of EAT-rv (p < 0.05).

Discussion

To our knowledge, this is the first large-scale clinical study to investigate the distribution of EAT and its association with metabolic syndrome. In this study, we explored a new measurement site for echocardiographic EAT thickness (EAT-ivg), which was greater than EAT-rv. The median values for EAT-rv and EAT-rv were 5.27 (4.01–6.86) and 3.01 (2.09–3.90), respectively. The EAT-rv values found among our sample of the Chinese population were similar to those of South Koreans, but lower than values measured among Caucasians. This racial difference was also confirmed by Fox et al. Our results show that EAT thickness was higher in older people and lower in males, a finding which is consistent with a study by Graeff et al. Both clinical studies and autopsies confirmed that EAT thickness significantly correlates with aging. As for gender difference, our results were consistent with the study by Baragetti et al., but opposite to the study by Calabuig et al. This discrepancy could be explained by the fact that the latter study did not include multivariate analysis.

In 2003, Iacobellis et al. first measured EAT thickness on the right ventricle anterior free wall (EAT-rv) with echocardiography, which showed an excellent agreement with MRI epicardial measurements and a significant association with metabolic syndrome. However, the association was reduced when the size of the study population was increased and multivariate analysis was performed. In our study, EAT-rv also lacked a significant correlation with metabolic syndrome after multivariate analysis, but EAT-ivg did significantly correlate with metabolic syndrome. In addition, the $\kappa$ value between EAT-rv and EAV was lower than that of EAT-ivg and EAV, and the AUC for the application of EAT-rv in predicting metabolic syndrome was lower than that of EAT-ivg. Therefore, EAT-ivg is a better index for echocardiographic EAT thickness and is more valuable in predicting metabolic syndrome.

We also analyzed the independent risk factors of EAT thickness. In addition to age and sex, waist circumference – a commonly used assessment tool for visceral fat – was also an independent risk factor. This conclusion is consistent with previous reports. High-density-lipoprotein cholesterol is a protective factor of EAT thickness, according to our results. In a cross-sectional study of 72 hemodialysis patients, a low HDL cholesterol level was identified as an independent risk factor of elevated EAT thickness. A meta-analysis showed that there was a highly significant (p < 0.001) correlation between EAT thickness and HDL cholesterol. Our results confirm the previous findings.

**Table 5. Association between EAT metrics and metabolic syndrome**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>EAT-rv</th>
<th>p-value</th>
<th>EAT-ivg</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Crude OR</td>
<td>1.337 (1.162–1.538)</td>
<td>&lt;0.001</td>
<td>1.628 (1.498–1.770)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Age-adjusted OR</td>
<td>1.306 (1.124–1.518)</td>
<td>&lt;0.001</td>
<td>1.569 (1.457–1.690)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Age- and waist-circumference-adjusted OR</td>
<td>1.163 (0.985–1.374)</td>
<td>0.075</td>
<td>1.363 (1.221–1.520)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Age-, waist-circumference- and BMI-adjusted OR</td>
<td>1.116 (0.938–1.329)</td>
<td>0.217</td>
<td>1.331 (1.182–1.498)</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

EAT – epicardial adipose tissue; EAT-rv – EAT thickness on the right ventricular anterior free wall; EAT-ivg – EAT thickness on the anterior interventricular groove; BMI – body mass index; OR – odds ratio.

Fig. 3. The AUC of EAT-rv and EAT-ivg applied in predicting metabolic syndrome.
Conclusions

In summary, EAT-ivg had a stronger agreement with EAV than EAT-rv, it was independently associated with metabolic syndrome and it was more valuable in predicting metabolic syndrome than EAT-rv. Therefore, the anterior interventricular groove can serve as a new measurement site which can better reflect EAT thickness.

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