Serum Adiponectin Levels are Associated with Microcirculatory Function, but not with Coronary Artery Disease in the Young

Dear Sir,

Adiponectin is secreted in mature adipocytes, has a role in diabetes and obesity and has been studied in coronary artery disease (CAD), considered cardioprotective,[1] what is nonetheless controversial.[2,3] The assessment of systemic microvascular reactivity is useful for the evaluation of cardiovascular diseases, and laser speckle contrast imaging (LSCI) is an innovative approach, using the cutaneous microcirculation as a representative vascular bed.[4] We evaluated serum adiponectin levels in early-onset CAD (EOCAD) patients and age-matched controls, as well as their association with microvascular function assessed by LSCI. EOCAD was defined as any obstruction ≥50% on coronary angiography, prior myocardial infarction, or myocardial revascularization in patients ≤45 years. Patients with acute coronary syndromes or interventions for <6 months were not included in the study. Controls were adults ≤45 years without CAD or cardiac symptoms. After a 12-h fast, venous blood was collected. Adiponectin was measured in serum by ELISA (Human High Molecular Weight Adiponectin, Millipore, Missouri, USA). Cutaneous microvascular reactivity was evaluated using LSCI (PeriCam; Perimed, Sweden), as previously described,[5] using transdermal iontophoretic delivery of acetylcholine (ACh) or sodium nitroprusside (NPS). Perfusion changes were measured in arbitrary perfusion units and expressed as peak values, representing the maximal vasodilation observed, and area under the curve of vasodilation. The study complied with the 1964 Declaration of Helsinki and its amendments and was approved by the local ethics committee. Informed written consent was obtained from all participants. Categorical variables were expressed as n (%) and compared with Chi-square. Continuous variables were expressed as a mean ± standard deviation or median/interquartile range, compared with Mann–Whitney’s test. Correlations were evaluated with Spearman’s test. A value of P < 0.05 was considered statistically significant.

Twenty-five EOCAD patients (age 42.8 ± 2.0 years, 60% male, 88% hypertensive, 28% diabetic, 56% obese) and 25 controls (41.1 ± 3.4 years, 44% male; 0, 4% and 24%, respectively) were studied. Patients had higher body mass index (BMI) (31.7 ± 5.8 vs. 26.2 ± 3.5 kg/m², P < 0.05), glucose (115.0 ± 31.0 vs. 93.5 ± 13.3 mg/dl, P < 0.05) and triglycerides (164.0 [46.0–52.0] vs. 91.0 [42.0–242.0] mg/dl, P < 0.05), and lower high-density lipoprotein (HDL)-cholesterol (36.6 ± 7.1 vs. 43.9 ± 10.2 mg/dl, P < 0.05). Serum adiponectin was not significantly different between patients and controls (19.7 [9.0–57.0] vs. 24.3 [4.0–59.0] µg/ml). There was a trend toward lower adiponectin levels in obese than in nonobese (18.1 [9–34] vs. 25.1 [4–59] µg/ml, P = 0.07). Significant correlations between adiponectin and other variables are shown in Table 1.

In conclusion, both endothelium-dependent and endothelium-independent vasodilatation were positively correlated with adiponectin levels, what reinforces the vasoprotective effect of adiponectin. The absence of difference of adiponectin levels among patients and controls may be attributed to the small sample size. To the best of our knowledge, this is the first study to use LSCI show the association between adiponectin and microcirculatory function. Further larger studies are needed to increase this knowledge.

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Conflicts of interest

There are no conflicts of interest.

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Table 1: Significant (P<0.05) correlations between serum adiponectin, clinical and microcirculatory variables

<table>
<thead>
<tr>
<th>Variable</th>
<th>Spearman’s coefficient</th>
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<tbody>
<tr>
<td>Body mass index</td>
<td>−0.38</td>
</tr>
<tr>
<td>HDL-cholesterol</td>
<td>0.41</td>
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<tr>
<td>ACh-induced peak vasodilatation (APU)</td>
<td>0.51</td>
</tr>
<tr>
<td>Area under the curve of ACh-induced vasodilatation (APU)</td>
<td>0.53</td>
</tr>
<tr>
<td>NPS-induced peak vasodilatation (APU)</td>
<td>0.61</td>
</tr>
<tr>
<td>Area under the curve of NPS-induced vasodilatation (APU)</td>
<td>0.56</td>
</tr>
</tbody>
</table>

ACh: Acetylcholine, APU: Arbitrary perfusion units, HDL: High-density lipoprotein, NPS: Sodium nitroprusside
References


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