Loss of the normal coupling between the anaerobic threshold and insulin sensitivity in chronic heart failure

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Abstract

Objective—To explore whether the anaerobic threshold, a measure of the balance between aerobic and anaerobic cellular metabolism, is related to whole body insulin sensitivity in healthy individuals and in patients with chronic heart failure, which involves an imbalance of aerobic and anaerobic metabolism.

Design—Case-control study.

Setting—A teaching hospital department specialising in heart failure.

Patients—20 healthy individuals (mean (SEM) age 55.2 (2.7) years) and 36 patients with chronic heart failure (59.1 (2.0) years, New York Heart Association class I–IV, anaerobic threshold 11.8 (0.7) ml/kg/min, left ventricular ejection fraction 26 (2)%).

Interventions—An intravenous glucose tolerance test for assessment of insulin sensitivity (minimal model analysis) and a maximum treadmill exercise test for assessment of the anaerobic threshold, derived from measurement of oxygen consumption and carbon dioxide output.

Main outcome measures—Relation between insulin sensitivity and the anaerobic threshold in patients with chronic heart failure.

Results—While anaerobic threshold was positively correlated with insulin sensitivity in healthy controls (r = 0.72, p < 0.001), no such relation was observed in patients with chronic heart failure. In stepwise multiple linear regression analyses of variables in healthy individuals, insulin sensitivity emerged as the only predictor of anaerobic threshold (standardised coefficient = 0.72, p < 0.001), while fasting insulin, incremental insulin area, and total body fat (dual photon x ray absorptiometry) failed to enter into final models (joint R = 0.52, p < 0.001).

Conclusions—In healthy individuals, whole body insulin sensitivity is related, or “coupled,” to the anaerobic threshold. The absence of such metabolic coupling in patients with chronic heart failure provides further evidence of disturbed cellular metabolism in patients with this condition.

Keywords: heart failure; anaerobic threshold; insulin resistance

The cellular supply of energy in the form of high energy phosphates relies on a balance between glycolysis (mainly anaerobic) and the citric acid cycle (aerobic). As well as requiring glucose and oxygen, the generation of glycolytic intermediates also requires insulin. This hormone not only drives cellular uptake of glucose, but also exerts a regulatory influence on glycolytic enzymes.1–3 Thus cellular metabolism requires coordination or “coupling” between the cellular processing of glucose and oxygen and insulin action.

On the basis of the above, we considered that whole body measures of the anaerobic threshold, which reflects the balance between aerobic and anaerobic metabolism, might relate to whole body insulin sensitivity. Our investigation focuses on healthy individuals and on patients with chronic heart failure—an insulin resistant state6 in which there is an imbalance between oxidative and non-oxidative metabolism.5–6

Methods

The study group consisted of 36 patients with chronic heart failure caused by coronary heart disease (19) or dilated cardiomyopathy (17). The diagnosis of chronic heart failure was based on standard criteria. All patients had been in heart failure for at least six months before the study. Concurrent treatment in the heart failure group included angiotensin converting enzyme inhibitors, loop diuretics, digoxin, warfarin, nitrates, amiodarone, and thiazide diuretics, either alone or in combination. No patients were taking hypouricaemic drugs. Healthy controls were matched for age and sex and were free from cardiovascular disease, on the basis of clinical history and examination and routine investigations. All patients gave written, informed consent and the study was approved by the local ethics committee.

Anaerobic threshold

During cardiopulmonary exercise testing, all patients were exercised to exhaustion (respiratory exchange ratio > 1.1). A one way valve connected to a respiratory spectrometer (Amin 2000, Odense, Denmark) was employed. Oxygen consumption was calculated on-line using a standard inert gas dilution technique.7 8 The slope of the regression line relating minute ventilation to carbon dioxide output was employed as an index of the ventilatory response to exercise.9 These measures were obtained during a maximum exercise test, using a modified Bruce protocol for patients with chronic heart failure.
The ventilatory anaerobic threshold was calculated according to the technique of Beaver et al.12 Briefly, the anaerobic threshold is selected as the oxygen consumption (VO2) at which the slope of the carbon dioxide output (VCO2) versus VO2 changes from 1 or slightly less than 1 to a slope which is steeper than 1. The intersection of these slopes is taken as the VO2 above which the increased VCO2 can only be explained by the increase in metabolic acidosis.

**INSULIN SENSITIVITY**

Participants underwent an intravenous glucose tolerance test with sampling at 16 time points during 180 minutes, as previously described.13 Dextrose was given intravenously as a 50% solution, at a dose of 0.5 g/kg body weight. Plasma glucose was determined on the same day using glucose oxidase procedures.14 Plasma insulin concentrations were measured on samples stored at −20°C using a radioimmunoassay procedure.15 Insulin sensitivity, inversely related to insulin resistance, was assessed using the minimal model approach.16

**BODY FAT COMPOSITION**

Height (m) and weight (kg) were determined to calculate body mass index (BMI: weight/height2). Total fat mass was estimated by dual energy x-ray absorptiometry (DXA) using a Lunar DPX (Lunar Corporation, Madison, Wisconsin, USA).17 All scans were performed and analysed using version 3.6z software. Precision of fat tissue measurements was better than 5%.18

**STATISTICAL ANALYSES**

These were carried out using the SYSTAT statistical package (SYSTAT Inc, Evanston, Illinois, USA). Univariate Pearson correlation coefficients were derived. A general linear model was used in multivariate analyses of pooled subjects. In all analyses, a p value of < 0.05 was considered statistically significant.

**Results**

As shown in table 1, the chronic heart failure group had a lower insulin sensitivity (p = 0.035) and anaerobic threshold (p < 0.001) as well as lower systolic (p = 0.002) and diastolic (p = 0.001) blood pressures.

Univariate analyses of variables in healthy individuals revealed a significant correlation between anaerobic threshold and insulin sensitivity (r = 0.72; p<0.001). The regression equation was AT = (8.95 + 4.3).Si and the coefficient of determination was R2 = 0.52. Univariate analyses of variables in heart failure patients revealed a significant correlation between anaerobic threshold and insulin sensitivity (r = −0.12; p = NS). The regression equation was AT = (5.57 + 3.1).Si and the coefficient of determination was R2 = 0.15.

**Table 1** Characteristics of the study and control groups

<table>
<thead>
<tr>
<th>Variable</th>
<th>Healthy controls (n = 20)</th>
<th>CHF patients (n = 36)</th>
<th>p Value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>55.2 (2.7)</td>
<td>59.1 (2.0)</td>
<td>NS</td>
</tr>
<tr>
<td>Body mass index (kg/m2)</td>
<td>28.2 (1.1)</td>
<td>25.6 (0.7)</td>
<td>NS</td>
</tr>
<tr>
<td>Total body fat (kg)</td>
<td>22.3 (1.8)</td>
<td>18.7 (1.2)</td>
<td>NS</td>
</tr>
<tr>
<td>LV ejection fraction (%)</td>
<td>ND</td>
<td>25.6 (2.3)</td>
<td></td>
</tr>
<tr>
<td>Systolic blood pressure (mm Hg)</td>
<td>126.0 (3.1)</td>
<td>111.8 (2.8)</td>
<td>0.002</td>
</tr>
<tr>
<td>Diastolic blood pressure (mm Hg)</td>
<td>79.4 (2.4)</td>
<td>70.0 (1.6)</td>
<td>0.001</td>
</tr>
<tr>
<td>NYHA class (n)</td>
<td>I – 4</td>
<td>II – 11</td>
<td></td>
</tr>
<tr>
<td></td>
<td>III – 18</td>
<td>IV – 3</td>
<td></td>
</tr>
<tr>
<td>Fasting glucose (mmol/l)</td>
<td>5.4 (0.1)</td>
<td>5.7 (0.3)</td>
<td>NS</td>
</tr>
<tr>
<td>Fasting insulin (pmol/l)</td>
<td>53.3 (~7.7, +9.0)</td>
<td>56.6 (~6.6, +7.5)</td>
<td>NS</td>
</tr>
<tr>
<td>Incremental glucose area (mmol/min)*</td>
<td>631.5 (147.1)</td>
<td>593.3 (35.1)</td>
<td>NS</td>
</tr>
<tr>
<td>Incremental insulin area (10^4.[pmol/l].min)**</td>
<td>2.52 (~0.25, +0.27)</td>
<td>2.02 (~0.27, +0.29)</td>
<td>NS</td>
</tr>
<tr>
<td>Insulin sensitivity (10^5/min/[pmol/l])‡</td>
<td>3.33 (~0.61, +0.67)</td>
<td>2.02 (~0.27, +0.29)</td>
<td>0.035</td>
</tr>
<tr>
<td>Anaerobic threshold (ml/kg/min)</td>
<td>16.8 (1.0)</td>
<td>11.8 (0.7)</td>
<td>&lt; 0.001</td>
</tr>
</tbody>
</table>

Values are mean (SEM) unless otherwise stated. *Differences between groups from analysis of variance; † log transformed for statistical analysis; ‡ square root transformed for statistical analysis.

CHF, chronic heart failure; LV, left ventricular; ND, not done; NYHA, New York Heart Association.

**Table 2** Univariate Pearson correlation analysis of anaerobic threshold against metabolic variables in healthy individuals and in patients with chronic heart failure (CHF)

<table>
<thead>
<tr>
<th>Variable</th>
<th>Controls (n = 20)</th>
<th>CHF (n = 36)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fasting glucose</td>
<td>−0.37</td>
<td>−0.08</td>
</tr>
<tr>
<td>Fasting insulin</td>
<td>−0.49*</td>
<td>−0.03</td>
</tr>
<tr>
<td>Incremental glucose area</td>
<td>0.18</td>
<td>−0.04</td>
</tr>
<tr>
<td>Incremental insulin area</td>
<td>−0.64***</td>
<td>0.23</td>
</tr>
<tr>
<td>Insulin sensitivity</td>
<td>0.72***</td>
<td>−0.12</td>
</tr>
<tr>
<td>Total body fat</td>
<td>−0.28</td>
<td>0.15</td>
</tr>
</tbody>
</table>

*p < 0.05; **p < 0.01; ***p < 0.001.

Figure 1 Regression scatterplots of insulin sensitivity (Si) against anaerobic threshold (AT) in (A) healthy individuals, and (B) patients with chronic heart failure. r, Pearson correlation coefficient.
sensitivity, incremental insulin area, and fasting insulin (table 2). In contrast, no significant correlations were observed in the chronic heart failure group (fig 1).

Variables which were found to be correlated in univariate analyses were entered in multivariate analyses. In stepwise multiple linear regression analyses of variables in healthy individuals, insulin sensitivity emerged as the only predictor of anaerobic threshold (standardised coefficient = 0.72, p < 0.001), while fasting insulin, incremental insulin area, and total body fat failed to enter into final models (joint R = 0.52, p < 0.001).

Discussion
We have shown that in healthy individuals, there is a strong relation between the anaerobic threshold and insulin sensitivity. This relation might be expected in view of the coordination that exists between oxygen and glucose uptake at the cellular level. A salient finding from this study is that the relation between the anaerobic threshold and insulin sensitivity observed in healthy individuals is absent in patients with chronic heart failure.

Our demonstration of a lower anaerobic threshold in patients with chronic heart failure is in keeping with the finding of an early switch to anaerobic metabolism in this condition, as is in keeping with the finding of an early switch threshold in patients with chronic heart failure, as exists in healthy individuals. A strong relation between oxygen and glucose uptake might be expected in view of the coordination that exists between oxygen and glucose uptake at the cellular level. On the basis of our findings, disturbances in cellular metabolism that occur in heart failure, such as a low anaerobic threshold, do not appear to be related to reductions in insulin sensitivity. Further studies are needed to determine which additional factors are responsible for the “uncoupling” of glycolysis and insulin mediated glucose uptake in patients with chronic heart failure.

We conclude that whereas in healthy individuals, the anaerobic threshold is positively related to insulin sensitivity, this relation is absent in patients with chronic heart failure. This observation is in keeping with the known mechanisms that link oxygen and glucose metabolism and insulin action at the cellular level. On the basis of our findings, disturbances in cellular metabolism that occur in heart failure, such as a low anaerobic threshold, do not appear to be related to reductions in insulin sensitivity. Further studies are needed to determine which additional factors are responsible for the “uncoupling” of glycolysis and insulin mediated glucose uptake in patients with chronic heart failure.

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Covered stent to treat co-existent coarctation and aneurysm of the aorta in a young man

A 23 year old man presented with hypertension and clinical, electrocardiographic, echocardiographic, and radiographic features of coarctation. Aortography delineated the coarctation (gradient 40 mm Hg) and a spherical aneurysm 3 cm across, originating at the coarctation. Under general anaesthesia, a cut-down and femoral arteriotomy was made and a 16 F sheath inserted. A super stiff Amplatz 0.035 inch guidewire was advanced across the coarctation. A 37.5 mm long (maximum diameter 22 mm) AneuRx (Medtronic, Watford, UK) self expanding, Nitinol mesh stent, covered with a stretchable polytetrafluoroethylene membrane, was placed across both the coarctation and the neck of the aneurysm under fluoroscopic guidance. Serial dilatation was made with 12 × 40 mm and 15 × 40 mm balloons (Cordis, Ascot, UK). Balloon dilatation alone was rejected because of the danger of rupture of the aneurysm. A conventional, uncovered stent might have “splinted open” a dissection plane or failed to seal off the aneurysm. With the covered stent, care had to be taken to avoid the left subclavian artery, thereby avoiding any danger of embolising the vertebral circulation. The flexible nature of this stent graft, in contrast to slotted tube designs, suited the curve of the aorta. The final appearance of the stented segment was smooth and patent, with a minimal residual pressure gradient and exclusion of the aneurysm. The temptation to pursue an angiographically wider lumen was resisted because the gradient had been largely abolished and because of the risk of graft rupture. Computed tomography 24 hours later confirmed persisting exclusion of the aneurysm. Six months later, the patient was well with a blood pressure of 120/80 mm Hg.