Determination of Maximal Oxygen Uptake Using Seismocardiography at Rest

Mikkel T Hansen1*, Birk M Grønfeldt1*, Tue Rømer1, Mathilde Fogelstrøm1, Kasper Sørensen2, Samuel E Schmidt2, Jørn W Helge1

1Department of Biomedical Sciences, University of Copenhagen, Copenhagen, Denmark
2Department of Health Science and Technology, Aalborg University, Aalborg, Denmark

Abstract

Introduction: Assessment of maximal oxygen consumption (\(\dot{V}O_2\)max) is an important clinical tool when examining both healthy and unhealthy populations, as a low \(\dot{V}O_2\)max is associated with cardiovascular disease and all-cause mortality. Aim: This study investigated the accuracy of a non-exercise test for assessment of \(\dot{V}O_2\)max using seismocardiography (SCG). Methods: 97 participants (20-45 years, 50 males) underwent a non-exercise test using SCG at rest in the supine position (SCG \(\dot{V}O_2\)max) and a graded exercise test to voluntary exhaustion on a cycle ergometer with indirect calorimetry (IC \(\dot{V}O_2\)max). An interim analysis was applied after 50 participants had completed testing (SCG \(\dot{V}O_2\)max 1.0) allowing for the algorithm to be modified (SCG \(\dot{V}O_2\)max 2.1). Results: SCG \(\dot{V}O_2\)max 2.1 (n=47, test set) estimation was 3.5 ± 1.8 mL·min\(^{-1}\)·kg\(^{-1}\) (p<0.001) lower compared to IC \(\dot{V}O_2\)max, with a Pearson correlation of \(r=0.65\) (p<0.0001) and a standard error of estimate of 7.1 mL·min\(^{-1}\)·kg\(^{-1}\). The coefficient of variation between tests was 8 ± 1%. Conclusion: The accuracy of \(\dot{V}O_2\)max assessment using SCG requires further optimization prior to clinical application, as SCG \(\dot{V}O_2\)max was systematically lower than IC \(\dot{V}O_2\)max, and only a moderate correlation together with considerable variation were observed between tests.

1. Introduction

Cardiorespiratory fitness (CRF) is positively associated with self-rated health [1] and inversely correlated with a high risk of cardiovascular disease (CVD) and all-cause mortality [2,3]. In 2016, the scientific statement from the American Heart Association [4] clearly recognised the importance of CRF assessment in clinical practice in order to improve patient management and CVD risk prediction. The gold standard method for quantifying CRF is a direct measurement of pulmonary gas exchange during a maximal exercise test for obtainment of maximal oxygen consumption (\(\dot{V}O_2\)max) [4]. Even though some of the proposed barriers with this method (cost associated with equipment and trained professionals) are becoming less problematic [5], it still requires a maximal effort from an individual, which is not always possible. The potential of non-exercise-based estimations are huge as it provides a rapid and inexpensive method of estimating \(\dot{V}O_2\)max in public health and clinical settings [4,6]. However, the validity of non-exercise tests are not yet satisfying [7,8]. In 2020, a clinical non-exercise method using seismocardiography (SCG) for estimation of \(\dot{V}O_2\)max was proposed [9]. SCG is a recording of the cardiac vibrations on the chest wall produced by the beating heart, with an accelerometer [10]. New advances in low-cost lightweight sensors, signal processing and machine learning [11] has made this technique suitable for optimization of non-exercise based estimations. Sørensen and colleagues showed that aortic valve closing (AC) information derived from the SCG signal was highly correlated with \(\dot{V}O_2\)max (r=0.80) and that a regression model using BMI, sex, age and mean AC peak to peak amplitude had a correlation of r=0.90 with the gold standard measurement of \(\dot{V}O_2\)max in a relatively small sample size (n=26) [9]. Therefore, the purpose of the current study was to investigate the accuracy of a clinical non-exercise test for assessment of \(\dot{V}O_2\)max using SCG in a larger cohort.

2. Methods

2.1. Participants

One hundred participants (50 males) aged between 18 and 45 years were included in the study. Data from 97 participants (50 males) are included, as three participants had invalid data regarding the non-exercise \(\dot{V}O_2\)max estimation. Exclusion criteria were current or previous cardiovascular disease, chronic medication, pregnancy or conditions that prevented maximal effort testing. Participants received both oral and written information about experimental procedures and possible risks associated with the study, before signing a written consent. The study was approved by the Science Ethical Committee of the greater region of Copenhagen, Denmark (H-17008748), and adhered to the principles of the Helsinki
Declaration. The study is prospectively registered at ClinicalTrials.gov (NCT03504306).

2.2. Study Design

An interim analysis was planned after the first 50 participants had completed testing (25 females / 25 males). This allowed for adjustment in the algorithm before a blinded analysis of the last 47 participant was performed. Hereby the first 50 participants were used as a training set and the remaining 47 as a test set.

The participants arrived at the laboratory after at least 4 hours fasting and without vigorous exercise performed within the last 24 hours before testing. The participants then underwent following measurements: Anthropometrics were measured and, after voiding and while wearing minimal clothing, body composition was determined by DXA scan (Lunar iDXA, GE Healthcare). Then, three measures of blood pressure (Boso-medicus control, Jungingen, Germany) were performed each separated by 2 minutes rest with an initial 5 minutes resting period. A 5 ml resting blood sample was hereafter obtained from the antecubital vein for assessment of HbA1c, haemoglobin, and haematocrit.

Table 1. Participant characteristics (n=97)

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Mean ± SD (Range)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, yrs.</td>
<td>31 ± 1 [20 - 45]</td>
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<tr>
<td>Height, cm</td>
<td>175 ± 2 [155 - 202]</td>
</tr>
<tr>
<td>Weight, kg</td>
<td>73.0 ± 2.3 [47.6 - 99.3]</td>
</tr>
<tr>
<td>BMI, kg·m⁻²</td>
<td>23.8 ± 0.5 [18.5 - 32.2]</td>
</tr>
<tr>
<td>Body fat, %</td>
<td>23.2 ± 1.4 [7.0 - 43.0]</td>
</tr>
<tr>
<td>Systolic BP, mmHg</td>
<td>125 ± 3 [92 - 165]</td>
</tr>
<tr>
<td>Diastolic BP, mmHg</td>
<td>76 ± 2 [56 - 96]</td>
</tr>
<tr>
<td>Resting HR, bpm</td>
<td>57 ± 2 [37 - 87]</td>
</tr>
<tr>
<td>Hemoglobin, mmol/L</td>
<td>8.6 ± 0.1 [7.2 - 10.3]</td>
</tr>
<tr>
<td>Hematocrit, %</td>
<td>41.2 ± 0.5 [35.0 - 49.0]</td>
</tr>
<tr>
<td>HbA1c, mmol/mol</td>
<td>32 ± 0.0 [25 - 39]</td>
</tr>
<tr>
<td>VO₂max, mL·min⁻¹·kg⁻¹</td>
<td>46.1 ± 1.4 [26.9 - 64.7]</td>
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Note: data are presented as mean ± 95% CI and [range]. BMI; body mass index; BP; blood pressure; HR; heart rate; HbA1c; glycated hemoglobin. b n=95.

A non-exercise VO₂max estimation using seismocardiography (SCG VO₂max) in the supine position following 5 minutes of bed rest was then conducted as previously described [9]. In brief, an ultra-sensitive accelerometer (Silicon Design 1521-002), with a resolution of ± 2 g, low noise at 7 µg / √ Hz and a frequency response 0-300 Hz, was placed on the lower part of sternum with double adhesive tape for recording of the SCG signal. The accelerometer measured 19 mm in width, 21 mm in length and 11 mm in height and weighed 5 grams including the electronic components and the ABS plastic housing. Resting ECG (a three lead ECG, with four electrodes placed on the right and left shoulder and right and left iliac crests) and SCG were recorded for 5 minutes using an iWorx IX-228/s (iWorx, Dover, New Hampshire) connected to a PC, acquisition unit sampling at 5000 Hz. LabScribe recording software (Version 3. Dover, New Hampshire) was used. Lastly, after a 5 minute warm-up at 75W participants performed a graded exercise test with 25W increments every minute until voluntary exhaustion on a cycle ergometer (Monark 839E, Monark Exercise AB). Pulmonary gas exchange measurements were obtained breath-by-breath during exercise and sampled in 10-s intervals by an automated online system (Quark CPET, COSMED). Gas analysers were calibrated with a compressed gas mixture (5% CO₂ and 16% O₂) and the digital flowmeter calibrated using a 3 L calibration syringe (COSMED) before each test. The VO₂max criteria was O₂ levelling off and a respiratory exchange ratio (RER) > 1.15.

2.3. Signal Processing

The ECG and SCG recordings were exported from the iWorx system and processed in MATLAB (2018a, MathWorks, Inc.) The signal processing was performed manually and this has previously been described [9]. The company behind the SCG VO₂max estimation model, VentriJect A/S, performed the signal processing and was blinded to the measured VO₂max, but did receive demographic data (weight, height and age) of the participants.

2.4. VO₂max Prediction Models

The features included in the two prediction models are:

SCG 1.0 VO₂max = 44.1 - 0.465 · BMI + 6.79 · SEX – 0.187 · AGE + 0.292 · ACpp

SCG 2.1 VO₂max = -65.895 + 0.06 · ACpp + 0.176 · tACp_p + 0.625 · FriendsAlgo – 155.4 · tIVCTRoustrr + 0.542 · S2FrequencySpec

Abbreviations: ACpp: peak to peak amplitude in SCG diastolic complex, tACp_p: time intervals between peaks in SCG diastolic complex, FriendsAlgo [6]; an algorithm based on sex, age and body weight for prediction of VO₂max, tIVCTRoustrr: robust estimation of isovolumetric contraction time normalised against the average duration of heart beats, S2FrequencySpec; frequency of the average SCG diastolic complex quantified using principal component analysis.

The SCG 1.0 refers to the previous model [9] and SCG 2.1 to the adjusted model. The SCG VO₂max 2.1 model was fitted to the data obtained previously [9] (n=43) and the initial 50 participants from this study, with a built-in function of MATLAB (stepwiselm) using both forward and backward stepwise regression. Independent variables were included or removed from the model based on statistical significance of the change in the sum of squared
errors. The criteria was p < 0.05 for addition and p < 0.10 for removal. This process was repeated until no more parameters could be added or removed.

### 2.5. Statistics

Data are presented as mean ± 95% confidence intervals (CI) with significance set at an α level of 0.05. Systematic difference in measured values between the initial 50 and the last 47 participants were analysed with an unpaired t test. Inter-method validity was analysed with a paired r test. Pearson correlation coefficient r, coefficient of variation (CV) and standard error of estimate (SEE) [9]. A Bland-Altman plot with 95% limits of agreement (LoA) assessed the agreement. Pearson correlation coefficients were interpreted as follows: very high >0.90, high 0.70-0.90, moderate 0.50-0.70, low 0.30-0.50 and little if any 0.00-0.30 [12]. Statistical analyses were performed, and figures constructed in GraphPad Prism 9.2.0 (Software Inc.) and Microsoft Excel (Microsoft Corporation).

### 3. Results

A systematic difference between the initial 50 participants and the last 47 was observed in the HbA1c measurement (33 ± 1 and 32 ± 1 mmol/mol, respectively) and IC VO₂max value (44.4 ± 1.6 and 47.9 ± 2.2 ml·min⁻¹·kg⁻¹, respectively) (p<0.05).

#### 3.1. Interim Analyses and Performance in Training Set

For SCG 1.0 a significant bias of -1.7 ± 1.5 ml·min⁻¹·kg⁻¹ with 95% LoA ranging ±10.3 ml·min⁻¹·kg⁻¹ was found, when compared to IC VO₂max (p=0.028). The correlation analysis revealed a correlation of r=0.60 (p<0.0001), with a SEE of 5.6 ml·min⁻¹·kg⁻¹. The intra-individual CV was 7 ± 1%. For SCG 2.1 a non-significant bias of -0.9 ± 1.3 ml·min⁻¹·kg⁻¹ with 95% LoA of 8.1 and -9.9 ml·min⁻¹·kg⁻¹ was found compared to IC VO₂max (p=0.172). The correlation was r=0.70 and with a SEE of 4.7 ml·min⁻¹·kg⁻¹. The CV between tests was 6 ± 1%.

#### 3.2. Test Set VO₂max Estimation

SCG VO₂max 2.1 was 7% lower (p<0.001) compared to IC VO₂max (Figure 1), with a moderate Pearson correlation observed between tests (Figure 2). Results from the accuracy assessment of the VO₂max estimation models, compared to gold standard IC VO₂max are presented in table 2.

![Figure 1. A Bland-Altman plot of the agreement between VO₂max estimated with a non-exercise model using seismocardiography (SCG 2.1) and directly measured with indirect calorimetry (IC).](image1.png)

![Figure 2. Scatterplot of the correlation between VO₂max estimated with a non-exercise model using seismocardiography (SCG 2.1) and directly measured with indirect calorimetry (IC).](image2.png)

### Table 2. Accuracy of the VO₂max estimation models compared with IC VO₂max (n=47, test set).

<table>
<thead>
<tr>
<th></th>
<th>FriendsAlgo</th>
<th>SCG 1.0</th>
<th>SCG 2.1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pearson, r</td>
<td>0.51</td>
<td>0.57</td>
<td>0.65</td>
</tr>
<tr>
<td>Bias, ml·min⁻¹·kg⁻¹</td>
<td>-6.4 ± 2.1</td>
<td>-4.2 ± 1.9</td>
<td>-3.5 ± 1.8</td>
</tr>
<tr>
<td>SEE, ml·min⁻¹·kg⁻¹</td>
<td>9.6</td>
<td>7.9</td>
<td>7.1</td>
</tr>
<tr>
<td>CV, %</td>
<td>12 ± 2</td>
<td>9 ± 2</td>
<td>9 ± 2</td>
</tr>
</tbody>
</table>

Note: data are presented as mean ± 95% CI. SEE; standard error of estimate, CV; coefficient of variation, IC; indirect calorimetry. FriendsAlgo [6].

### 4. Discussion

The accuracy of VO₂max determination using seismocardiography at rest was investigated in the present study. The interim analysis of the first 50 participants using the SCG 1.0 model revealed beside a significant bias, a moderate correlation of r=0.60 compared with IC VO₂max, which is lower compared to results previously obtained...
with that model (high, r=0.90) [9]. The SEE was also higher in the present study (5.6 vs. 3.2 ml·min⁻¹·kg⁻¹, respectively) [9]. The participants included in the present study constituted a broader representative of the normal population in regards to VO₂max values as the participants were solely in the low to moderate VO₂max category in the previous study [9]. VO₂max estimation accuracy in the training set was improved in the modified SCG 2.1 model, as no significant bias, a high correlation and smaller variations were found between tests. When applied in the test set, the SCG 2.1 revealed only a moderate correlation along with a significant bias and larger variations between tests. In addition, the Bland-Altman plot shows a negative proportional bias (Figure 2), which when combined with a significantly higher measured VO₂max between the initial 50 and the last 47 participants, reduces the accuracy and thus makes the SCG 2.1 model less accurate when estimated in a more representative extract of the population with higher VO₂max values. However, when the SCG 2.1 test set is compared with the FriendsAlgo estimation (which includes a training set of 7783 subjects and validation set of 1287 subjects), the accuracy was higher as both systematic bias, correlation and variations were improved (Table 2). Nevertheless, further acquisitions of VO₂max data from a broad population are required in order to improve accuracy of the prediction model.

5. Conclusion

The SCG signal contains information which can improve estimation of VO₂max, however the accuracy of VO₂max assessment using SCG requires further optimization prior to clinical application, as SCG 2.1 VO₂max was systematically lower than the gold standard measurement with indirect calorimetry, and only a moderate correlation and considerable variations were observed between tests. Ongoing development of the prediction model is in progress in order to improve the accuracy of VO₂max estimation using SCG.

Acknowledgments and Author Contributions

We would like to thank all the participants for their commitment to the study and VentriJect A/S for the equipment lent and data analyses regarding SCG VO₂max estimation. JWH and BMG designed the study. BMG and MF carried out data acquisition. MTH, BMG, TR, KS, SES and JWH contributed to data analyses and interpretation. MTH wrote the manuscript. All authors revised and approved the manuscript. *denotes shared first authorship.

Conflict of Interest

The study was financially supported by VentriJect A/S with no restriction on publication. SES and KS holds significant shares in VentriJect A/S and works part time as Chief Scientific Officer and Head of Software in VentriJect A/S, respectively.

References


Address for correspondence:
Mikkel Thunestved Hansen
Blegdamsvej 3, DK-2200 Copenhagen, Denmark
Mikkel.hansen@sund.ku.dk