Validity of the 6 min walk test in prediction of the anaerobic threshold before major non-cardiac surgery

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The authors conclude that those walking >563 m do not require CPET, and those walking <427 m do.

Patients who walk a distance between the two cut-off points need further evaluation.

The findings of this study provide important validation of simple walk test in risk stratification and prognosis.

Editor’s key points

- The 6 min walk test was compared with cardiopulmonary exercise testing (CPET) in predicting anaerobic threshold.
- The authors conclude that those walking >563 m do not require CPET, and those walking <427 m do.
- Patients who walk a distance between the two cut-off points need further evaluation.
- The findings of this study provide important validation of simple walk test in risk stratification and prognosis.

Background. For perioperative risk stratification, a robust, practical test could be used where cardiopulmonary exercise testing (CPET) is unavailable. The aim of this study was to assess the utility of the 6 min walk test (6MWT) distance to discriminate between low and high anaerobic threshold (AT) in patients awaiting major non-cardiac surgery.

Methods. In 110 participants, we obtained oxygen consumption at the AT from CPET and recorded the distance walked (in m) during a 6MWT. Receiver operating characteristic (ROC) curve analysis was used to derive two different cut-points for 6MWT distance in predicting an AT of ≤11 ml O2 kg⁻¹ min⁻¹; one using the highest sum of sensitivity and specificity (conventional method) and the other adopting a 2:1 weighting in favour of sensitivity. In addition, using a novel linear regression-based technique, we obtained lower and upper cut-points for 6MWT distance that are predictive of an AT that is likely to be (P≤0.75) <11 or >11 ml O2 kg⁻¹ min⁻¹.

Results. The ROC curve analysis revealed an area under the curve of 0.85 (95% confidence interval, 0.77–0.91). The optimum cut-points were <440 m (conventional method) and <502 m (sensitivity-weighted approach). The regression-based lower and upper 6MWT distance cut-points were <427 and >563 m, respectively.

Conclusions. Patients walking >563 m in the 6MWT do not routinely require CPET; those walking <427 m should be referred for further evaluation. In situations of ‘clinical uncertainty’ (≥427 but ≤563 m), the number of clinical risk factors and magnitude of surgery should be incorporated into the decision-making process. The 6MWT is a useful clinical tool to screen and risk stratify patients in departments where CPET is unavailable.

Keywords: anaerobic threshold; exercise test; oxygen consumption; preoperative care

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Service infrastructure costs may prohibit setting up a CPET service. Subjective functional assessment of METs, although a simpler alternative, has been shown to have user and physiological limitations. An alternative, simple, objective measure of exercise capacity may therefore more robustly aid risk stratification, where CPET is unavailable. Ideally, such a test should be validated against measured CPET parameters.

A review of the validity data supporting functional exercise tests revealed the 6 min walk test (6MWT) to be the most extensively researched and established test for use in clinical or research contexts in the cardiorespiratory domain. Previous studies have demonstrated a positive correlation between CPET measurements and distance walked in patients with cardiorespiratory disease. Although the 6MWT has been shown to predict outcome after pulmonary resection and lung volume reduction surgery, there is no literature pertaining to major non-cardiac surgery. We believe that based on this evidence and pilot data from...
our institution, the 6MWT might be suitable to provide the simple, objective assessment of exercise capacity outlined above.

The aim of this study was to assess the validity of the distance walked during the 6MWT in predicting the AT (and other parameters) derived from CPET.

Methods

The protocol for this concurrent validity study was approved by the National Research and Ethics Service in August 2008 (08/H1305/62). Trial registration: ISRCTN 12656789.

Participants were recruited from the preoperative assessment clinics at the James Cook University Hospital between October 2008 and January 2010. After verbal explanation and a patient information sheet, written informed consent was obtained.

Participants included in the study were aged 50–85 yr and awaiting scheduled major non-cardiac surgery (Grade 3 or 4 surgery as defined by NICE guidance). 19 Exclusion criteria comprised: medical contraindication to CPET20 or failure to complete a baseline CPET, lower limb claudication and inability to maintain a steady walking pace on level ground. After a medical screening examination, patients were invited to participate.

For a desired precision of estimation of ±0.10 (95% confidence interval width) around a postulated validity correlation coefficient of $r=0.70$ (for 6MWT distance in the prediction of AT) derived from pilot work, a sample size of 100 patients was estimated. Allowing for an attrition rate of 25%, a final sample size of 125 participants was required. A total of 186 individuals were screened for inclusion. Of these, 129 participants were enrolled. Characteristics, co-morbid diseases, surgical procedures undertaken, and medications prescribed for participants completing both CPET and 6MWT (119 participants) are shown in Supplementary Table S1.

Participants were asked to complete two exercise tests: CPET (on a cycle ergometer) and a 6MWT. The CPET was performed first, in order to screen for significant cardiovascular pathology, thus ensuring the safe conduct of the 6MWT. To minimize participant inconvenience, both tests were undertaken on the same day. After CPET, patients were provided with refreshments and allowed an appropriate rest interval between tests. The 6MWT was only undertaken once the participants had reported that they had no residual fatigue from CPET. To avoid study bias, the 6MWT was administered by an investigator blinded to the results of the CPET.

Cardiopulmonary exercise test

The CPET was performed using the Medgraphics Ultima system (Tewkesbury, Gloucestershire, UK) and a Lode Corival V2 cycle ergometer (BV Medical Technology, Groningen, The Netherlands). Flow and gas calibrations were performed before each test session, which was subsequently conducted to our standard protocol (available in Supplementary material). All usual patient medication was continued.

The test was terminated when the participant reached volitional exhaustion ($V_{O2}$ peak) or earlier if another termination criterion was fulfilled. The V-slope comparison plot was compiled using Breeze software (Medgraphics) and interpreted by two trained observers on completion of all study testing (G.R.D. and R.C.F.S.).

Six min walk test

After successful completion of CPET, participants performed the 6MWT as outlined in the guidance published by the American Thoracic Society (ATS). 19 Individuals walked to their own maximum pace along a flat corridor, marked with a 30 m track, aiming to cover as much distance as possible in the timed 6 min. Participants wore a MIR0xi pulse oximeter (Medical International Research, Roma, Italy) to record heart rate response and oxygen saturations.

The ATS suggest that a practice test is not needed in most settings. 19 Furthermore, data from our pilot study (unpublished observation) confirmed that the test was highly reproducible, with an intraclass correlation coefficient (ICC 3.1) of 0.94, and a non-substantial mean bias of 18 m greater on a second walk. Thus, a single 6MWT was performed in the current study.

Test outcome measures recorded

- CPET—oxygen consumption at the AT (using the V-slope technique), 23 oxygen consumption at volitional exhaustion ($V_{O2}$ peak), the $VE/ V_{CO2}$ recorded at AT, and maximum heart rate achieved ($HR_{max}$)
- 6MWT—maximal distance walked and $HR_{max}$

Statistical analysis

Ordinary least-squares linear regression models were applied to obtain the validity coefficient ($r$) and the standard error of the estimate (SEE)—the typical error associated with the prediction of AT (or $V_{O2}$ peak or $VE/ V_{CO2}$ slope) from 6MWT distance in an individual patient. Receiver operating characteristic (ROC) curve analysis was used to derive cut-points for 6MWT distance for the prediction of AT ($<11 ml \ O2 kg^{-1} min^{-1}$, $AT <8 ml \ O2 kg^{-1} min^{-1}$, $V_{O2}$ peak $<15 ml \ O2 kg^{-1} min^{-1}$, and a combination of AT $<11 ml \ O2 kg^{-1}$ min$^{-1}$ and $VE/ V_{CO2}$ slope $>34$. The optimum cut-point was determined as the value corresponding with the greatest accuracy (highest sum of sensitivity plus specificity; i.e. with sensitivity and specificity weighted equally). When a test is to be used for screening purposes and risk stratification, however, a cut-off value with greater sensitivity (fewer false-negatives) may be desirable. Therefore, we derived an alternative cut-point by adopting a 2:1 weighting for sensitivity:specificity.

To refine the ROC-derived cut-offs, we used the obtained regression equation and SEE, to derive lower and upper cut-points for 6MWT distance that are predictive of an AT that is likely to be less than or greater than these prognostic AT thresholds. (A 6MWT distance falling between these two cut-points is assumed to be in an area of ‘clinical uncertainty’.)
Herein, ‘likely to be’ is defined as a probability of ≥0.75 (odds of at least 3:1 in favour) of the patient’s true AT being less than (lower cut-point for 6MWT distance) or greater than (upper cut-point) 11 ml O2 kg⁻¹ min⁻¹, given the predicted AT from the regression equation and the observed prediction error (SEE). This probability is derived from the disposition of the confidence interval for the predicted value to the prognostic value of 11 ml O2 kg⁻¹ min⁻¹ and is calculated using the Student t-distribution. The required t-value is derived as the prognostic value for AT minus the predicted value from the regression and divided by the obtained SEE: (11–9.7)/1.9=0.68. The area under the t-distribution to the left of this value with the appropriate degrees of freedom is 0.75, providing the probability that the patient’s true AT is <11 ml O2 kg⁻¹ min⁻¹ if their predicted value from the regression was 9.7 ml O2 kg⁻¹ min⁻¹. Rearranging the derived linear regression equation gives the 6MWT distance predictive of an AT of 9.7 ml O2 kg⁻¹ min⁻¹; this is the lower cut-point. The upper cut-point was calculated in an identical fashion. All analyses were conducted using StatsDirect (Altrincham, UK; v. 2.7.8) and Medcalc (Mariakerke, Belgium; v. 11.5) software packages.

We adopted an objective criterion to identify and remove outliers with a standardized residual of >3.6 from the analysis. With the assumption of normality, this threshold identifies values that would occur only rarely (<5% of the time) with this sample size.

Results

In total 119 of 129 recruited participants completed both exercise tests. Of the 10 individuals not completing: two withdrew consent after CPET, one failed to reach AT during CPET, five were unable to complete the full 6 min of walking, and two individuals had no reason documented. Of the 119 participants, an additional seven participants were eliminated from the analysis due to a persistently elevated respiratory exchange ratio (RER) likely a consequence of hyperventilation due to anxiety, poor accommodation to the mouthpiece, or both. In such cases, an AT is still detectable but it will be a ‘pseudo-threshold’ occurring before the actual AT resulting in an underestimation.

Screening for severe outliers resulted in the removal of one case for the AT analysis and one case for the V̇O₂ peak analysis, resulting in a data set of n=110 complete cases.

Exercise test results

The CPET and 6MWT results for study participants are presented in Table 1. The peak exercise challenge was comparable between the two tests as judged by the similar mean maximum heart rate. Figure 1 illustrates a scatter plot of AT vs 6MWT distance.

Linear regression analyses to predict the AT, V̇O₂ peak, and V̇E/V̇CO₂ from the distance walked during the 6MWT are shown in Table 2, and the results of the ROC curve analyses are detailed in Table 3. The area under the ROC curve indicates that a randomly selected individual from the positive group (AT <11 ml O2 kg⁻¹ min⁻¹) has a 6MWT distance value (Y) smaller than that of a randomly chosen individual from the negative group (X) 85.2% of the time [P(Y<X)=0.852]. The likelihood ratios indicate that a 6MWT distance of ≤440 m is obtained around 15 times as frequently in patients with an AT of <11 ml O2 kg⁻¹ min⁻¹ than in those with an AT above this threshold and that a 6MWT distance of ≥440 m is obtained approximately a third as frequently in patients with an AT of <11 ml O2 kg⁻¹ min⁻¹ than in those with an AT above this value. The ROC curve for this analysis is shown in Figure 2, illustrating that the area under the curve is substantially larger than that of 'no

| Table 1 Exercise test results [mean (SD)]. HRmax, maximum heart rate achieved; AT, anaerobic threshold; V̇O₂ peak, peak oxygen consumption |
|----------------------|----------------------|----------------------|----------------------|
| Outcome              | Slope               | Intercept            | Correlation coefficient (r) (95% CI) | Standard error of the estimate (SEE) (95% CI) |
| CPET                 | 0.019               | 1.598                | 0.68 (0.56–0.77)         | 1.9 (1.7–2.2) ml O2 kg⁻¹ min⁻¹ |
| AT                   | 0.033               | −0.326               | 0.75 (0.65–0.82)         | 2.7 (2.4–3.1) ml O2 kg⁻¹ min⁻¹ |
| V̇O₂ peak             | −0.028              | 48.479               | 0.46 (0.30–0.60)         | 5.0 (4.4–5.8) |

**Table 2 Linear regression analyses with 6MWT distance as the predictor. AT, anaerobic threshold; V̇E/V̇CO₂, ventilatory equivalents for carbon dioxide; V̇O₂ peak, peak oxygen consumption**
Discrimination' (0.50) indicated by the diagonal. Figure 3 shows the dot plot for 6MWT distance in the two groups.

Lower and upper cut-points for 6MWT distance derived from regression analysis

From the regression modelling, the lower and upper cut-points for 6MWT distance predictive of a true AT that is likely to be \( (P \geq 0.75) \) <11 or >11 ml O\(_2\) kg\(^{-1}\) min\(^{-1}\), respectively, were below 427 m (positive test) or above 563 m (negative test). For the 8 ml O\(_2\) kg\(^{-1}\) threshold for the AT, the lower and upper cut-points were <269 and >405 m. For the V\(_O2\) peak <15 ml O\(_2\) kg\(^{-1}\) min\(^{-1}\), the lower and upper cut-points were <409 and >520 m.

Discussion

In this study, we have confirmed that the 6MWT may be a useful practical method for risk stratification, with a large effect size observed for the correlation between the distance walked during a 6MWT and oxygen consumption at both AT and V\(_O2\) peak. AT, measured during CPET, is presently recognized as the most robust endpoint to inform perioperative risk stratification.\(^4\) –\(^7\) For this reason, the majority of our analysis and inference focuses on the relationship between 6MWT distance and AT, rather than V\(_O2\) peak.

The ROC curve analyses for both the <11 and <8 ml O\(_2\) kg\(^{-1}\) min\(^{-1}\) thresholds for AT revealed that the 6MWT

Table 3 ROC curve analyses. AT, anaerobic threshold; VE/\(V\_CO2\), ventilatory equivalents for carbon dioxide; V\(_O2\) peak, peak oxygen consumption

<table>
<thead>
<tr>
<th>CPET measurement (ml O(_2) kg(^{-1}) min(^{-1}))</th>
<th>AT &lt;11</th>
<th>AT &lt;8</th>
<th>AT &lt;11 and VE/(V_CO2) &gt;34</th>
<th>V(_O2) peak &lt;15</th>
</tr>
</thead>
<tbody>
<tr>
<td>ROC curve sensitivity: specificity weighting</td>
<td>1:1</td>
<td>2:1</td>
<td>1:1</td>
<td>2:1</td>
</tr>
<tr>
<td>Prevalence (%)</td>
<td>58.2</td>
<td>19.1</td>
<td>36.4</td>
<td>51.8</td>
</tr>
<tr>
<td>AUC</td>
<td>0.852</td>
<td>0.857</td>
<td>0.801</td>
<td>0.856</td>
</tr>
<tr>
<td>95% CI</td>
<td>0.771–0.912</td>
<td>0.778–0.917</td>
<td>0.741–0.871</td>
<td>0.776–0.916</td>
</tr>
<tr>
<td>Cut-point (m)</td>
<td>&lt;440</td>
<td>&lt;502</td>
<td>&lt;411</td>
<td>&lt;450</td>
</tr>
<tr>
<td>Sensitivity</td>
<td>0.641</td>
<td>0.844</td>
<td>0.857</td>
<td>0.952</td>
</tr>
<tr>
<td>95% CI</td>
<td>0.511–0.757</td>
<td>0.731–0.922</td>
<td>0.637–0.970</td>
<td>0.762–0.999</td>
</tr>
<tr>
<td>Specificity</td>
<td>0.957</td>
<td>0.674</td>
<td>0.843</td>
<td>0.697</td>
</tr>
<tr>
<td>95% CI</td>
<td>0.852–0.995</td>
<td>0.520–0.805</td>
<td>0.750–0.911</td>
<td>0.590–0.790</td>
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<td>Positive likelihood ratio</td>
<td>14.73</td>
<td>2.59</td>
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<tr>
<td>95% CI</td>
<td>12.1–17.9</td>
<td>2.1–3.2</td>
<td>4.5–6.6</td>
<td>2.7–3.7</td>
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<tr>
<td>Negative likelihood ratio</td>
<td>0.38</td>
<td>0.23</td>
<td>0.17</td>
<td>0.07</td>
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<tr>
<td>95% CI</td>
<td>0.09–1.5</td>
<td>0.1–0.5</td>
<td>0.05–0.5</td>
<td>0.01–0.5</td>
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</table>

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distance is an adequate discriminator between high and low AT patient groups. However, deriving an optimum single cut-point from ROC curve analysis is challenging and not ideally suited to a clinical context. Sensitivity can be weighted to reduce the false-negative rate, but we believe that the regression-based analysis represents a refinement of single ROC curve cut-points allowing for clinical variation and uncertainty. Using the regression method, a patient with a positive test (6MWT < 427 m) is likely to be at high perioperative risk, and a patient with a negative test (6MWT > 563 m) would be considered low risk. A patient completing a distance of ≥ 427 but ≤ 563 m is in a zone that we define as ‘clinical uncertainty’. We can usefully incorporate these regression analysis-derived risk categories into clinical practice (see below).

The current international guidance relies on subjective assessment of functional capacity, in the form of METs, as one of the three key variables in the decision-making process of risk stratification before non-cardiac surgery. A functional capacity of < 4 METs (inability to climb a flight of stairs) represents the threshold to trigger the high-risk limb of the risk stratification pathways. In the current study only one of 101 individuals (1%) reported a functional capacity of < 4 METs (14 ml O2 kg min⁻¹), whereas 58.2% of our participants had an objectively measured AT of < 11 ml O2 kg min⁻¹ during CPET. Interestingly, the individual reporting a functional capacity of < 4 METs attained an AT and VO₂ peak of 11.8 and 15.2 ml O2 kg min⁻¹, respectively, thereby representing low risk based on objective testing. We believe that based on our data, the 6MWT represents a superior and more robust technique for risk stratification than a self-reported cut-point of < 4 METs.

The other two key variables utilized to determine preoperative risk in the current international guidance are the number of clinical risk factors and nature of surgical intervention. Utilizing this approach in combination with 6MWT distance could help identify the most at risk individuals before surgery. The major benefits would be the ease with which it could be administered, minimal staff training and equipment requirements, and simple and quick to perform. In addition, the test is repeatable, is safe to perform, and entails a minimal increase in patient attendance time. Although not recommended as a replacement for CPET, the 6MWT could, in effect, act as a surrogate ‘sieve’ in identifying high-risk individuals who may require further assessment or optimization before surgery. In hospitals where CPET would perhaps not be utilized frequently, the 6MWT could be used as a cheap accurate alternative enabling identification and referral for CPET via loco-regional preoperative networked arrangements.

We believe that the upper and lower cut-points derived from the regression analysis provide the ideal platform in providing for such a model. For example, no further assessment would be required in an individual walking > 563 m during the 6MWT (upper cut-point, true AT likely to be > 11 ml O2 kg min⁻¹), whereas a patient walking < 427 m (lower cut-point, true AT likely to be < 11 ml O2 kg min⁻¹) would be considered high risk and should be referred for further functional assessment. In individuals walking a distance in the area of ‘clinical uncertainty’ (≥ 427 but ≤ 563 m), it would be important to incorporate the number of clinical risk factors and magnitude of surgical intervention into this clinical decision-making process, before consideration of further investigation. Therefore, a patient walking, say, 500 m together with two to three clinical risk factors should be further assessed, whereas an individual walking the same distance with a good health profile would not.

Our study is unique in being the first to examine the use of the 6MWT before operation in patients undergoing non-cardio-thoracic surgery. However, we identified three studies within cardio-respiratory medicine reporting similar correlations between 6MWT distance and CPET measurements to ours. These studies predominantly concentrated on correlations between peak oxygen consumption and 6MWT distance, reporting validity coefficients from r = 0.64 to 0.88. The observed correlations between 6MWT distance and AT and VO₂ peak in the current study are substantially larger than those reported in patients with chronic obstructive pulmonary disease.

Our results appear to conflict with those reported from a study of the validity of an intermittent shuttle walk test in assessing fitness for surgery, with the authors concluding that the discriminatory ability of the test was poor. However, a robust comparison of our findings with this study is not possible, as the patient group was substantially fitter (mean AT 12.7 vs 10.2 ml O2 kg min⁻¹) than our sample, and the authors do not detail the method used to determine the single ROC cut-points, nor the sensitivity, specificity, and likelihood ratios associated with the derived cut-points.

It is important to acknowledge a number of limitations to our study. First, we are utilizing a specific cut-off value for AT to discriminate between high- and low-risk individuals (a threshold value of 11 ml O2 kg min⁻¹). However, this threshold remains robust, despite being unchanged since proposed originally. Indeed, Snowden and colleagues reported a very similar AT cut-point (10.1 ml O₂ kg min⁻¹) in prediction of increased postoperative morbidity. Similarly, Wilson and colleagues reported that an AT of <11 ml O₂ kg min⁻¹ was a clinically significant predictor of mortality in major non-cardiac surgery patients.

Second, using the 6MWT as a surrogate provides limited diagnostic information on cardiorespiratory reserve, which can be obtained with CPET. We are however in effect suggesting the 6MWT as an improvement over a subjective cut-point of < 4 METs in identifying high-risk individuals and not in replacement of CPET. With the current financial constraints on the National Health Service in the UK, we believe that the 6MWT represents a robust pragmatic improvement where CPET is unavailable. Indeed, identification of high-risk individuals utilizing the 6MWT may enable streamlined pathways of care at the loco-regional level as outlined above. Such a tertiary referral service would be more cost-effective and avoid unnecessary...
duplication of tests. Third, the 6MWT is of limited utility in assessing patients with limb ischaemia or major limitation to exercise, for example, lower limb arthritis. Patients who cannot walk at a good pace have a resultant decreased 6MWT distance when compared with exercise results during non-weight-bearing cycle exercise. Finally, this study was designed to examine the prediction of CPET parameters from a 6MWT and not powered to predict perioperative outcome. We acknowledge that this might be considered an important direction for future research were the 6MWT to be adopted into regular clinical practice.

In conclusion, our results demonstrate that the 6MWT can be used robustly at preoperative assessment to assess exercise capacity. Where CPET is unavailable, we believe the regression analysis model presented provides an accurate, simple, and cheap way of clinically guiding further patient management as part of a preoperative screening process.

Supplementary material
Supplementary material is available at British Journal of Anaesthesia online.

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Declaration of interest
None declared.

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