

Treadmill Measured vs. Questionnaire Estimated Changes in Walking Ability in Patients With Peripheral Artery Disease

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- 1 Treadmill-measured versus questionnaire-estimated changes in walking ability in patients
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- 3
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18 WHAT THIS STUDY ADDS

In patients with peripheral artery disease, the Walking Estimated-Limitation Calculated by 19 History (WELCH) questionnaire score correlates with the treadmill maximal walking time. The 20 21 changes in WELCH score tend to decrease more than the objective changes in walking 22 impairment, particularly with a longer a test-retest interval. After revascularisation, a short-lived "honeymoon" (overestimation of the objectively measured change on the treadmill) is observed. 23 24 For long test-retest intervals, self-reported worsening according to the WELCH score should probably be confirmed on the treadmill before a decision to revascularise is taken. Whether other 25 questionnaires estimating walking impairment face the same issue remains to be determined. 26

28 ABSTRACT

Introduction: Determining the maximal walking time (MWT) using the treadmill test is the gold-standard method for evaluating walking capacity and treatment effect in patients with peripheral arterial disease (PAD). However, self-reported functional disability is important when assessing the quality of life. We compared changes in the Walking Estimated Limitation Calculated by History (WELCH) questionnaire scores with the MWT.

Methods: A transversal study was performed among patients with intermittent claudication. The 34 treadmill test (3.2 km/h; 10% grade) and WELCH questionnaire were administered to all patients 35 for objective evaluation of walking capacity. Given the log-normal distribution of these 36 37 parameters in PAD patients, a log transformation was applied to the WELCH score (LnW) and maximal walking time (LnT). The responsiveness of the WELCH score was determined using 38 39 mean changes and correlation coefficients of LnW and LnT changes. The effect of time on the "Estimated minus Real" (E-R) changes (LnW-change minus LnT-change) was assessed after 40 categorization of patients into various test-retest intervals. Patients who underwent lower-limb 41 42 revascularisation between the two tests and those who underwent medical treatment only were 43 analysed.

Results: Correlation coefficients between LnW and LnT for tests 1 and 2 were r = 0.514 and r =
0.503, respectively (P < 0.001, for both). Correlation for LnW-change vs. LnT-change was 0.384
(P < 0.001). E-R was positive only early after surgery. E-R was negative for all test–retest
intervals >1 year in revascularised and non-revascularised patients.

48 Conclusions: Changes in WELCH scores correlated with changes observed on the treadmill in
49 patients with intermittent claudication. For long test-retest intervals, WELCH changes tended to

50 overestimate the worsening of walking impairment as compared with the measured difference

observed in both revascularised and non-revascularised patients. A short-lived "honeymoon"

- 52 (overestimation of the benefit for the shortest test-retest interval) was observed only in
- 53 revascularised patients.
- 54 KEYWORDS: Peripheral artery disease; Walking impairment; Treadmill test; Questionnaire;
 55 Revascularisation

57 INTRODUCTION

Estimation of walking impairment through standard questionnaires is easy, can be routinely 58 performed and scored, and is of major interest in patients with peripheral artery disease (PAD) 59 and claudication.¹ Questionnaire scores are generally validated against different objective 60 measurements of walking ability.²⁴ The effect of time between the two evaluations, specifically 61 62 from the lower-limb revascularisation procedure upon the concordance of subjective to objective changes, has not been analysed. The "walking estimated limitation calculated by history" 63 (WELCH) questionnaire⁵ is a relatively simple tool to self-report walking limitations in PAD. It 64 can be self-administered and compares favourably with previously proposed available tools.^{2,6} It 65 is currently available in different languages and is easy to use in routine practice.⁷⁻⁹ Correlation of 66 between the WELCH score and objective measures of walking capacity ranges from 0.58 to 67 $0.82^{6,8,9}$ and it is not impaired by age.² To date, the reliability and sensitivity to changes of the 68 WELCH have not been studied extensively. The effect of the test-retest interval on WELCH 69 changes compared with the changes in treadmill measured maximal walking time (MWT) is 70 71 unknown.

This study determined whether the WELCH questionnaire was sensitive to changes in walking capacity and how the WELCH score changes compared with objectively measured MWT changes. Next, we aimed to determine whether the relationship between WELCH and MWT changes varied with the test-retest interval and if there was a difference between the two tests in patients that did not undergo revascularisation between the two estimates (medical treatment only) and in those subjected to arterial lower-limb revascularisation.

78 MATERIALS AND METHODS

A transversal study was performed among patients referred to our laboratory for walking 79 test investigations. Most patients when primarily referred to us complained of claudication for 80 months and most had previously benefited from optimal medical treatment for PAD. Retest visits 81 82 were either based on the evaluation of residual limb or non-limb symptoms or sometimes on the surgeon's request for an objective evaluation of functional improvement. Eligibility included: 83 age>18 years, the ability to walk on a treadmill, the ability to understand the study goals and 84 85 instructions of the tests, and self-completed questionnaires. The study was approved by the 86 Institutional Review Board and was registered on CNIL (Commission Nationale Informatique et 87 Liberté). It was performed according to the International Ethics Standards and conforms to the Helsinki Declaration. Patients were aware that the results were being recorded during treadmill 88 tests and that this investigation could be used for research purposes and were informed of their 89 right to oppose. All investigators participating in the study were informed on how to perform the 90 investigations, complete the files, and score the questionnaires. The study was registered with 91 92 ClinicalTrials.gov Identifier: NCT01424020.

93 Completion of questionnaires

Each patient was provided a pen and reading glasses (if needed) and received oral instructions for 94 completing the questionnaire on arrival at the laboratory. Each patient self-completed the 95 96 questionnaires while in the waiting room of the laboratory before the walking test was performed. 97 The questionnaire included: date of the visit, history and on-going treatments, name and surname, age, sex, body weight, stature, active smoking, and the WELCH in French. The WELCH is a 98 99 four-item questionnaire that can be self-administered. The original questionnaire was developed in French but is currently available in various languages. In brief, the first three items refer to the 100 maximal time that can be sustained when walking at different walking speeds. Answers to each 101

item include proposals ranging from impossible (zero points) to 3 hours or more (7 points). The 102 103 fourth and last item requires the patient to estimate his/her usual walking speed as compared to that of his/her siblings or of people of comparable age. Possible answers ranged from "much 104 105 slower" (coefficient = 1) to "faster" (coefficient = 5). The WELCH score was calculated as follows. One is subtracted from the sum of the points of the three first items. The result of this 106 107 subtraction was multiplied by the coefficient of item four. Assuming that patients can walk at least 30 seconds at a low speed, the final score ranges from zero (severe disability) to 100 (ability 108 109 to walk faster than other people for at least 3 hours). Note that patients were never reminded of 110 their answers to the previously completed WELCH questionnaires or their previous walking distance on the treadmill. A typical example of filling of the WELCH questionnaire is provided 111 112 in Fig. 1.

113 Clinical data and investigations

114 Patients were admitted to the test room and their usual walking speed was measured between two 115 lines drawn on the floor separated by 10 meters, which were traced on the floor between the waiting room and the testing room. As in our previous studies, patients unable to walk 10 m in 116 less than 15 seconds were considered unable to walk 3.2 km/h on the treadmill and underwent a 117 specific protocol on the treadmill;¹⁰ they were excluded from the study. Technicians, nurses, or 118 119 physicians supervised the completion of the questionnaires and completed any eventual 120 incomplete items or clinical characteristics from the patient's most recent file or by immediate measurement. We systematically recorded the WELCH score, presence/absence of lower limb 121 122 revascularisation (bypass surgery or angioplasty) in the past or since the first visit for returning patients, age, sex, body weight, stature, and the ankle to brachial systolic pressure index (ABI). 123

Last, we retrieved the type of revascularization for the revascularised patients from their medicalfile or from contact with the surgeon (from patients referred from private practice physicians).

126

127 Treadmill test

To assess walking capacity, all patients performed a standardised constant load treadmill test 128 under medical supervision, at a constant speed of 3.2 km/h, at a 10% incline progressively 129 reached in 1 minute. For patients who were able to walk 15 minutes (900 sec) at a constant 130 workload, at minute 15, the protocol was changed to an incremental load protocol with steps of 1-131 minute duration until exhaustion or pain limitation.¹¹ The treadmill tests were performed by 12-132 133 lead ECG monitoring. The test variable used during the treadmill test was the MWT, defined as the time that severe claudication pain forced cessation of exercise or as the time that the test was 134 interrupted for medical reasons (severe cardiac arrhythmia, abnormal repolarization, etc.). 135

136 Data analysis

From the laboratory database, patients who had undergone at least two different consecutive tests 137 were selected. For patients who had performed multiple tests, only the last two visits were 138 analysed because of data accessibility. We previously reported that in patients with limiting 139 claudication both MWTs treadmill constant load tests¹² and WELCH scores⁶ show a log-normal 140 distribution in PAD patients complaining of exertional limb pain. Subsequently, the MWT and 141 WELCH scores were log-transformed for analysis and were referred to as LnT and LnW, 142 respectively. Changes in MWT and WELCH scores were calculated as the difference in the LnT 143 144 or LnW determined in the second test from the respective LnT and LnW determined at the first

visit and were analysed for the whole population. From previous studies, we estimated the 145 Spearman "r" coefficients of the correlation between LnT and LnW changes to be .30. This was 146 determined from previous studies, which showed the correlation of the walking impairment 147 questionnaire score to maximal walking distance was .33,⁴ and because the correlation of the 148 WELCH score and treadmill MWT was found to range from .58–.61;⁶ thus, the resulting 149 estimation of the correlation to be expected for WELCH and MWT changes was the square of the 150 "r" correlation values: .34–.37. For the alpha two-tailed .05 and 80% power, the minimal 151 152 observation number was 85.

153 Thereafter, patients were divided into two groups: patients who underwent medical treatment only between their two tests (non-revascularised group) and patients who had some form of lower 154 155 limb revascularisation between their two tests (S group). In each group, time intervals were categorized into six test-retest intervals as follows: test-retest intervals ranging from 0 to <6 m, 6 156 m to <12 m, 1 to <2 y, 2 to <3 y, 3 to <4 y, and 4 y or more. For each test-retest interval, the 157 median duration of the test-retest interval was recorded. Within each interval, the mean and SEM 158 of the difference between LnW-changes and LnT-changes was calculated. This difference was 159 160 noted as E-R. In practice, E-R was assumed to reflect overestimation of the benefit or 161 underestimation of the impairment (positive difference) or inverse underestimation of the benefit or overestimation of the impairment (negative difference) of the subjective estimation of walking 162 impairment evolution (WELCH questionnaire) as compared to the "real" objective evolution in 163 the MWT between the two visits. We performed a database analysis, on the observation of at 164 least six non-revascularised and revascularised patients, in each test-retest interval. 165

166 Statistical analysis

Results are presented as mean±SEM when normally distributed, as median [25°–75° centiles] 167 when appropriate, and as percentages. The unpaired t-test, Mann-Whitney test, and Chi-squared 168 test were used to compare non-revascularised and revascularised patients. ANOVA with 169 170 Dunnett's post-hoc test was used to compare patients within different test-retest intervals with patients with the shortest (0 to <6 m) interval. Two-tailed paired t-tests were used for LnT and 171 LnW to compare results of the first and second test, respectively. Correlation of the LnW-172 changes to LnT-changes was analysed for the whole population. Non-linear logarithmic models 173 174 were chosen as apparent models that best fitted the values observed for mean E-R for the 6 test-175 retest intervals in non-revascularised and revascularised patients, respectively. Statistical analyses were performed using SPSS V15.0 (SPSS Inc. LEADTOOLS®, LEAD Technology Inc.). For all 176 177 tests, a two-tailed p-value <.05 was used to indicate statistical significance.

178

179 **RESULTS**

This study included 346 consecutive patients with a mean age of 61.9 years; 87% were men. A 180 181 flowchart of recruitment is presented in Fig. 2. Baseline characteristics of the study population 182 are presented in Table 1. Among revascularised patients, 52 had an aortic and/or iliac angioplasty, 31 femoral and/or popliteal angioplasty, 14 aortic and/or iliac bypass surgery, and 12 183 a femoral and/or popliteal bypass surgery between their two tests. No significant difference was 184 found between non-revascularised and revascularised patients relative to morphology or 185 186 treatments. However, at baseline, both PAD (lower ABI) and walking impairment were more 187 severe (lower WELCH score and lower maximal walking time on treadmill) in revascularised 188 than in non-revascularised patients. The WELSH scores of the 346 patients were 20 (10-33) vs.

189 22 (10-39) Wilcoxon P = 0.27, based on the first and second test, respectively. Table 2 reports the
190 major characteristics of the revascularised and non-revascularised patients as a function of test191 retest interval.

No complications were noted during the walking test, although 153 (44.2%) and 149 (43.1%) of patients reported dyspnoea, fatigue, or chest pain during tests one and two, respectively. Fiftynine of the patients studied at test 1 and 94 of the patients studied on test 2 reached the incremental phase of the treadmill test (MWT >900 s).

The correlation coefficient between LnW and LnT was r = .514 in test 1 and r = .503 in test 2, respectively (p <.01 for both) (Fig. 3). The correlation for LnW-change vs. LnT-change was .384 (p <.01).

The LnW and LnT values observed for the six test-retest intervals are shown in Fig. 4. On 199 200 average, the revascularised patients showed an increase in walking time (LnT was 0.65 for the shortest Test-retest interval and 0.58 for the largest test-retest interval) after revascularisation, 201 while the non-revascularised patients had minimal, if any, increase in walking time between their 202 203 tests, irrelevant of the test-retest interval (LnT being slightly above or close to zero). Of note, in 204 most patients, the first evaluation was performed while medical treatment was optimal and not when the diagnosis of PAD was determined. Therefore, non-revascularised patients generally 205 presented stable, previously diagnosed, claudication. Only 18 patients were evaluated during the 206 207 first test before the onset of medical therapy (start of antiplatelet and/or cholesterol lowering drugs) or before referral to a rehabilitation program. Furthermore, the change in self-reported 208 evaluation of walking capacity (through the change in WELCH score) decreased with an increase 209 in the test-retest interval. LnT started from 0.99 and decreased to -0.02 in revascularised patients 210

211 and decreased from 0.01 to -0.53 in non-revascularised patients. As a result, the E-R difference 212 (Fig. 5) was negative in all except one case and roughly decreased with an increase in the test-213 retest intervals. The only exception was for the shortest interval corresponding to the pre- and 214 early post-revascularisation estimation in the non-revascularised group. This was the only group where the subjective estimation of changes by the patients (LnW of the Welch score) was, on 215 216 average, higher than the objective measurement of changes (LnT of the MWT on the treadmill). Note that the SEM of the 2–3-year interval was large given the limited number of observations (n 217 218 = 6).

219

220 DISCUSSION

The present study aimed to determine whether the WELCH questionnaire is sensitive to changes 221 in walking capacity and how the WELCH score changes compared with objectively measured 222 changes maximal walking time. We show that the WELCH score is as consistent in objective 223 224 measurements as other more complex tools and is an easy-to-use instrument that can determine changes in the walking capacity of patients with PAD. The correlation coefficient that we 225 226 identified (r = .38) may appear low; however, it is higher than the objective and subjective instruments of other studies. A previous study that assessed the effect of an intervention and 227 observed a larger range of change found a correlation of changes in the treadmill distance with 228 changes in the distance sub-scores of the walking impairment questionnaire ranging from r = .31229 to r = .34.^{13, 14} Comparable results were reported with Short-form-36 physical summary score 230 changes of r = .29 or using the intermittent claudication questionnaire changes of 0.38.^{13, 14} 231

The original observation of the effect of interval duration on the relationship between WELCH 232 and MWT changes in non-revascularised and patients is of specific importance. The non-233 revascularised patients showed no major differences in walking capacity (change in LnT close to 234 235 zero) for the various time intervals, while changes in LnT were >0 for all test-retest intervals in non-revascularised patients. While the shortest test-retest LnW interval was also close to zero, 236 237 LnW decreased with longer test-retest intervals. Thus, the E-R decreased with the increase in the test-retest interval. We previously demonstrated that the WELCH was independent of age and our 238 239 largest interval was only five years. Depression and anxiety are frequent symptoms in cardiovascular patients.¹⁵⁻¹⁷ Whether mood changes could explain our results for long test-retest 240 intervals remains to be determined. Overall, determining whether self-reported or treadmill-241 measured changes should prevail in the decision to revascularise a PAD patient is an open debate. 242 Another interesting observation is the initial transient overestimation (positive E-R) in 243 revascularised patients. Conversely, "overestimation of the benefit" in revascularised patients 244 could result from fear of a negative outcome before arterial revascularisation.¹⁸ However, it is 245 246 well known that initial excellent results one year after lower limb arterial revascularisation progressively worsen with time.¹⁹ Subsequently, many patients become symptomatic again on a 247 248 mid-term and long-term basis after arterial revascularisation, facing the same issues of chronic 249 pain and disease as non-revascularised patients. Nonetheless, the positive E-R phenomenon is a short-lived honeymoon. 250

Determining the changes in walking ability and evolution of walking impairment in patients with PAD is of major importance in clinical routine trials and research trials. Laboratory investigations (among which treadmill testing is the recommended "gold standard"), are time consuming, and are not necessarily accessible to physicians in routine clinical settings. Furthermore, results are 255 influenced by the protocols used and may not optimally reflect overall walking impairment. 256 Although subjective, questionnaires are of interest when accounting for the perceived impact of PAD in a patient's life. Various disease-specific or non-disease-specific tools are available. The 257 258 "walking impairment questionnaire", the "intermittent claudication questionnaire", the "claudication scale", and the « vascular quality of life" questionnaires are generally considered 259 260 specific for patients with PAD.^{13, 20-22} These questionnaires are long and impractical, with each including >14 items. A shorter version of the original 25-item the « vascular quality of life" is 261 262 available, but it focuses on the impact of walking impairment on usual activities, rather than on the severity of walking impairment itself.²³ Among these tools, the walking impairment 263 questionnaire is the most widely used tool with >150 references from a Medline search. 264 Unfortunately, the questionnaire is lengthy, subject to errors when self-completed, and cannot be 265 scored simply by mental calculations.^{24, 25} 266

The fact that studied groups included different patients may represent an issue, with small clinical 267 differences between the groups. A prospective study with repeated measurements would be a 268 269 better approach; nevertheless, the feasibility of a prospective approach over a 4-year period is tentative due to technical and financial reasons. In this transversal observational study, no 270 271 predefined visits were proposed according to a predefined interval to the patients. Thus, there 272 may be a bias with patients referred to us for multiple tests that are not comparable to general 273 PAD patients. A second issue involves the interval between tests that may slightly differ from the interval between revascularisation and the second visit in the revascularised group. This is only 274 true for the largest test-retest interval because for the shortest intervals, the first visit generally 275 276 preceded the revascularisation by approximately a few weeks. A third limitation is that we did not account for co-morbid conditions that may have occurred (or become exacerbated) between 277

the interval of the two tests because the data submitted for analysis focused only on vascular 278 279 diseases. It cannot be excluded that the self-reported limitation could be influenced by an underlying cause (vascular vs. non-vascular) of the walking impairment. To date, the WELCH 280 281 survey has not been tested in non-vascular populations. Obviously, the difference in self-reported and measured changes observed with test-retest interval in the revascularised group could also 282 depend on a deterioration of the revascularisation with time in successful vs. unsuccessful 283 revascularisation procedures. Unfortunately, we did not have access to primary patency of 284 285 revascularisation. Of interest is the fact that the MWT difference between test 2 and test 1 286 remained positive for revascularised patients, while the WELCH score difference was indeed positive for short test-retest intervals but decreased with larger intervals (Fig. 4). Furthermore, the 287 time evolution in the self-reported vs. measured changes observed in the non-revascularised 288 patients (Fig. 5) was similar to that of the revascularised patients, despite apparently comparable 289 290 ABIs for the non-revascularised groups. Thus, it appears that the evolutions of differences in 291 changes are unrelated to haemodynamic changes

292

293 CONCLUSION

The WELCH questionnaire is a valid tool to detect changes in the daily walking ability of patients with intermittent claudication. The self-reported WELCH score tends to decrease more than an objective measurement of walking impairment, especially if the test-retest interval is long. After revascularisation, a short-lived "honeymoon" (overestimation of the objectively measured change on the treadmill) may be observed. This honeymoon appears to last <1 year,

| 299 | after which revascularised patients follow the same evolution as non-revascularised patients for |
|-----|--|
| 300 | large test-retest intervals. |

- 301 In PAD patients, a self-reported worsening identified using the WELCH score during two
- 302 consecutive visits >1 year apart should probably be systematically confirmed on a treadmill,
- 303 before a decision to revascularise is taken. Whether other questionnaires aiming at estimating
- 304 walking impairment face the same issue remains to be determined.

305

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TABLES

Table 1. Baseline characteristics of the study population

| Study population | No revascularisation | Revascularisation | P-value |
|----------------------------|----------------------|-------------------|---------|
| Number of subjects | 237 | 109 | - |
| Age, mean (SD) | 61.6 (9.3) | 62.5 (9.6) | .400 |
| Men, % | 86.9 | 87.3 | .952 |
| Body mass index, mean (SD) | 27.1 (4.4) | 27.1 (4.6) | .891 |

| Antiplatelet drugs | 88.3 | 89.8 | .638 |
|---|---------------|---------------|------|
| Antihypertensive drugs | 75.1 | 69.3 | .293 |
| Cholesterol-lowering drugs | 79.9 | 81.6 | .679 |
| Anti-diabetic agents | 24.7 | 26.6 | .734 |
| History of previous lower limb bypass | 41.2 | 32.0 | .101 |
| Resting ankle-brachial index, mean (SD) | 0.80 (0.22) | 0.73 (0.21) | .004 |
| Current smoker, % | 34.9 | 36.6 | .762 |
| WELCH | 22 [12-36] | 18 [10-28] | .005 |
| LnW | 2.98 (0.84) | 2.72 (0.84) | .005 |
| Maximal walking time (s) | 324 [209-756] | 213 [150-345] | .001 |
| LnT | 5.90 (0.81) | 5.51 (0.66) | .001 |
| | | | 1 |

418 Results are mean (SD: Standard deviation) or Median [25-75 centiles] or percentages; WELCH =

419 Walking estimated limitation calculated by history; LnW = logarithmic value of WELCH score;

420 LnT = logarithmic value of walking time on treadmill

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Table 2. Major characteristics of patients in each of the test-retest intervals of non-revascularised
(medically treated) and revascularised patients. Ankle-brachial index (ABI) is the value at first

425 visit

| | Test-retest interval | Number of subjects | Age, mean | Men, % | BMI (kg/m²) | Ankle-brachial index | ABI change | Current smoker |
|------------------------------|----------------------|--------------------|-------------|--------|-------------|-------------------------|--------------|----------------|
| | 0 to <6 m | 47 | 60.0 (9.9) | 74.5 | 26.2 (4.2) | 0.78 (0.24) | 0.01 (0.18) | 36.2 |
| ion | 6 to <12 m | 57 | 63.3 (8.5) | 78.9 | 27.9 (4.6) | 0.82 (0.20) | -0.01 (0.19) | 22.9 |
| No revascularisation | 1 to <2 y | 69 | 62.4 (8.9) | 92.7 | 27.6 (4.2) | 0.80 (0.24) | -0.04 (0.19) | 27.5 |
| evascu | 2 to <3 y | 35 | 61.8 (10.7) | 94.2 | 26.3 (4.7) | 0.84 (0.19) | -0.05 (0.22) | 14.3 |
| No r | 3 to <4 y | 19 | 60.6 (8.5) | 100 | 26.6 (3.8) | 0.82 (0.17) | 0.01 (0.15) | 31.6 |
| | 4 y or more | 10 | 56.7 (10.1) | 100 | 26.8 (4.7) | 0.68 (0.15) | -0.01 (0.14) | 10.0 |
| E | 0 to <6 m | 34 | 60.5 (10.7) | 76.4 | 26.6 (4.1) | 0.68 (0.17) | 0.19 (0.22) | 32.4 |
| Lower limb revascularisation | 6 to <12 m | 33 | 64.0 (8.5) | 90.1 | 27.4 (4.8) | 0.75 (0.20) | 0.17 (0.27) | 21.2 |
| ascula | 1 to < 2 y | 18 | 63.4 (10.0) | 100 | 25.8 (4.7) | 0.70 (0.28) | 0.03 (0.21) | 38.9 |
| nb rev | 2 to < 3 y | 6 | 59.7 (8.8) | 100 | 27.9 (4.3) | 0.87 (0.24) | 0.07 (0.26) | 16.7 |
| wer lin | 3 to < 4 y | 10 | 65.9 (8.1) | 100 | 26.3 (3.1) | 0.74 (0.16) | 0.11 (0.12) | 30.0 |
| Lo | 4 y or more | 8 | 64.5 (11.3) | 100 | 30.2 (6.4) | 0.83 (0.32) | 0.03 (0.23) | 12.5 |

- 426 *Results are expressed as mean (standard deviation (SD)) or percentage; BMI, body mass index;*
- 427 no significant change for any variable from the first group on non-revascularised and
- 428 *revascularised patients except for sex.*

429

430 FIGURE LEGENDS

431 Fig. 1: English version of the WELCH questionnaire with example of scoring

432 Fig. 2: Flowchart of database analysis and resulting study population

433 Fig. 3: Scatterplot of logarithmic transformed changes in WELCH score (LnW) and maximal

434 walking time (LnT) in the first (test 1) and second (test 2) tests, in patients who received medical

435 treatment (non-revascularised) or underwent revascularisation (revascularised) between tests one436 and two.

437 **Fig. 4:** Mean logarithmic transformed WELCH score (LnW) and maximal walking time (LnT)

438 observed for the six test-retest intervals in patients who received medical treatment (non-

439 revascularised : non-S) or revascularisation (S) between tests one and two.

Fig. 5: Changes (test 2-test 1) in logarithmic transformed WELCH scores (LnW) and maximal walking time (LnT) observed for the six test-retest intervals. *R* is the Spearman coefficient of correlation of the regression analysis of average LnW-changes and LnT changes. The grey zone is the "honeymoon period" during which the self-reported difference was superior to the measured difference in the treadmill test for patients with lower limb revascularisation. Each point indicates the mean and standard error of mean. Please answer each of the following 4 items by placing an "X" in the box that best describes your situation. Please mark only one box per item. If you never perform an activity, estimate what it would be like if you did perform it. For the first 3 items, if you think that you would not be able to perform a specified task for at least 30 seconds without stopping to rest, please answer "impossible".

For each of the three following activities, how long can you perform the task easily on level ground & without stopping when ...

1/ ... walking slowly (slower than usual speed of relatives, friends, or other people of your own age)?

| Impossible | 30 seconds | 1 minute | 3 minutes | 10 minutes | 30 minutes | 1 hour | 3 hours |
|------------|------------|----------|-----------|------------|------------|--------|---------|
| | | | | | | | or more |
| | | | | | | | |

2/ ... walking normally (same as usual speed of relatives, friends, or other people of your own age)?

| Impossible | 30 seconds | 1 minute | 3 minutes | 10 minutes | 30 minutes | 1 hour | 3 hours or more |
|------------|------------|----------|-----------|------------|------------|--------|--------------------|
| | | | | | | | |
| 0 | 1 | 2 | 3 | 4 | . 5 | 6 | 7 |

3/ ... walking quickly (faster than usual speed of relatives, friends, or other people of your own age)?

| Impossible | 30 seconds | 1 minute | 3 minutes | 10 minutes | 30 minutes | 1 hour | 3 hours |
|------------|------------|----------|-----------|------------|------------|--------|---------|
| | | | | | | | or more |
| X | | | | | | | |
| 0 | 1 | 2 | 3 | 4 | 5 | 6 | 7 |

Compared to the usual walking speed of your relatives, friends, or people of your own age, do you think that you personally usually walk ... (Tick only one box)

much slower moderately slower a bit slower at the same speed faster



THANK YOU: You should have 1 box per item ticked... please check.

<u>WELCH score</u> = $[(5 + 2 + 0) - 1] \cdot 3 = 18$









