Feasibility of cardiopulmonary exercise testing and training using a robotics-assisted tilt table in dependent-ambulatory stroke patients

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Abstract

Background: We evaluated the feasibility of an augmented robotics-assisted tilt table (RATT) for incremental cardiopulmonary exercise testing (CPET) and exercise training in dependent-ambulatory stroke patients.

Methods: Stroke patients (Functional Ambulation Category ≤ 3) underwent familiarization, an incremental exercise test (IET) and a constant load test (CLT) on separate days. A RATT equipped with force sensors in the thigh cuffs, a work rate estimation algorithm and real-time visual feedback to guide the exercise work rate was used. Feasibility assessment considered technical feasibility, patient tolerability, and cardiopulmonary responsiveness.

Results: Eight patients (4 female) aged 58.3 ± 9.2 years (mean ± SD) were recruited and all completed the study. For IETs, peak oxygen uptake (V'O₂peak), peak heart rate (HRpeak) and peak work rate (WRpeak) were 11.9 ± 4.0 ml/kg/min (45 % of predicted V'O₂max), 117 ± 32 beats/min (72 % of predicted HRmax) and 22.5 ± 13.0 W, respectively. Peak ratings of perceived exertion (RPE) were on the range "hard" to "very hard". All 8 patients reached their limit of functional capacity in terms of either their cardiopulmonary or neuromuscular performance. A ventilatory threshold (VT) was identified in 7 patients and a respiratory compensation point (RCP) in 6 patients: mean V'O₂ at VT and RCP was 8.9 and 10.7 ml/kg/min, respectively, which represent 75 % (VT) and 85 % (RCP) of mean V'O₂peak. Incremental CPET provided sufficient information to satisfy the responsiveness criteria and identification of key outcomes in all 8 patients.

For CLTs, mean steady-state V'O₂ was 6.9 ml/kg/min (49 % of V'O₂ reserve), mean HR was 90 beats/min (56 % of HRmax), RPEs were > 2, and all patients maintained the active work rate for 10 min: these values meet recommended intensity levels for bouts of training.

Conclusions: The augmented RATT is deemed feasible for incremental cardiopulmonary exercise testing and exercise training in dependent-ambulatory stroke patients: the approach was found to be technically implementable, acceptable to the patients, and it showed substantial cardiopulmonary responsiveness. This work has clinical implications for patients with severe disability who otherwise are not able to be tested.

Keywords: Cardiopulmonary fitness, Exercise testing, Exercise training, Peak oxygen uptake, Robotics, Robotics-assisted tilt table, Stroke

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Background

Cardiopulmonary fitness is compromised in stroke patients: their peak oxygen uptake \((V’O_{2peak})\) ranges from 8–22 mL/kg/min, which corresponds to approximately half of age and gender matched healthy controls [1, 2]. The low \(V’O_{2peak}\) limits patients’ ability to live independently [3] and hinders participation in rehabilitation and exercise programmes [4]. Low cardiopulmonary fitness can further heighten the existing risk for cardiovascular disease [5] by predisposing patients to a sedentary lifestyle because of activity limitation and early fatigue [6].

A recent joint statement from the American Heart Association and the American Stroke Association recommends that stroke patients should undergo cardiopulmonary exercise testing (CPET) [4]. CPET delivers objective measures which allow accurate quantification of cardiorespiratory fitness, delineation of the physiological systems underlying exercise responses, and identification of exercise-limiting pathophysiological mechanisms [7]. CPET outcomes can also be used to evaluate the effects of a longitudinal training programme and to determine the training intensity for individualized exercise prescription [4, 8, 9]. However, impairments following stroke such as weakness, ataxia or spasticity can preclude some patients from exercise testing on standard devices. Semi-recumbent cycle ergometers and total-body recumbent steppers have hitherto been used as alternatives to standard treadmills and cycle ergometers in order to test patients with balance and coordination problems [4].

Despite increasing availability of adapted devices, suitable methods and data for patients who are severely disabled are lacking. This problem is clearly demonstrated in the systematic review by Smith et al. [1], where only 2 of the 42 studies included reported data from dependent-ambulatory patients. The authors pointed out that the exclusion of the severely disabled group may result in overestimation of cardiopulmonary fitness for the entire stroke population [1]. Another systematic review on the effects of cardiovascular exercise early after stroke pointed out that concepts to influence and evaluate cardiopulmonary fitness in severely disabled patients are still lacking [10]. These open questions are addressed in the present work.

A robotics-assisted tilt table (RATT) is a device used clinically for early rehabilitation in severely impaired and bedridden neurological patients. It tilts the patient upright, provides support with a body harness, promotes weight bearing on the feet and moves the legs in a cyclic stepping movement. To promote active participation during the rehabilitation process, we have augmented a RATT system to allow patients to see their exercise work rate together with a target work rate [11]. This approach was shown to be feasible for exercise testing both in normal subjects and in spinal cord injured patients [12, 13]. We hypothesized that the augmented RATT should enable stroke patients with severe motor weakness to be tested.

Incremental CPET aims to approach a person’s limit of functional capacity in regard to cardiopulmonary and/or neuromuscular exertion. The main parameters investigated here, which can be determined from incremental CPET, include both peak and submaximal values:

- **Peak:** peak oxygen uptake \((V’O_{2peak})\), which represents aerobics capacity; peak heart rate \((HR_{peak})\); and peak work rate \((WR_{peak})\), which is the highest volitional effort. Additional criteria are applied to \(V’O_2\) and \(WR\) responses to determine whether the observed peaks represent true maximal values.
- **Submaximal:** the 1st ventilatory threshold, denoted here as VT, which provides an approximation of endurance capacity [7, 14]; and the 2nd ventilatory threshold, denoted here as the respiratory compensation point (RCP), that occurs at the onset of hyperventilation [15].
- **Subjective measures** such as rating of perceived exertion (RPE) may also be recorded at intervals throughout the test.

This is the first study where the novel augmented RATT system was applied to severely-disabled patients. Therefore we wanted to investigate whether, with this new exercise testing modality, the principal incremental CPET outcomes can be identified and whether, during constant-load CPET, sustained exercise intensity meets recommendations for training.

The aim of this study was therefore to evaluate the feasibility of the augmented RATT for incremental cardiopulmonary exercise testing and exercise training in dependent-ambulatory stroke patients, i.e. those with severe physical disability who are unable to use standard devices. Criteria for the feasibility assessment were: (i) implementation – technical feasibility of the augmented RATT for exercise testing, (ii) acceptability – was the exercise tolerable?, and (iii) responsiveness – was there a measurable, high-level cardiopulmonary reaction?.

Materials and methods

Study design and participants

This descriptive, cross-sectional feasibility study was conducted at the Reha Rheinfelden, a rehabilitation centre in the north-west of Switzerland, from October 2013 to April 2014. The Ethics Review Committee of Canton Aargau, Switzerland, approved the study. All subjects gave their written informed consent before participating in the study.

Eight patients (4 female) aged 58.3 ± 9.2 years (mean ± SD) were recruited and all completed the study. The mean Functional Ambulation Category (FAC, [16]) was 1.8 (range 0–3; Table 1). Patient inclusion criteria were: (1) a
diagnosis of first-ever stroke, either ischaemic or intracerebral haemorrhage by radiologic evidence; (2) > 2 months post stroke; (3) > 18 years old; (4) dependent in ambulation with Functional Ambulation Category (FAC) ≤ 3; (5) Mini Mental State Examination (MMSE, [17]) score > 20 (cognitive function); and (6) willing to cooperate in the study and able to attend all testing sessions. Exclusion criteria were: (1) any contraindications to maximal exercise testing according to the American College of Sports Medicine guidelines [18]; (2) any contraindications for the RATT based on guidelines from the manufacturer; (3) severe aphasia or other communication problems; and (4) severe concurrent pulmonary disease. A cardiologist reviewed all prospective subjects for cardiac status before giving approval for formal enrolment.

### Table 1 Characteristics and demographic data of subjects (n = 8)

| Characteristic                    | Value                        |
|-----------------------------------|------------------------------|
| Age (years)                       | 58.3 ± 9.2                   |
| Sex, n (%)                        | Male 4 (50 %)                |
|                                  | Female 4 (50 %)              |
| Height (cm)                       | 167.6 ± 8.8                  |
| Body mass (kg)                    | 75.2 ± 7.4                   |
| Body mass index (kg/m²)           | 26.9 ± 3.3                   |
| Type of stroke, n (%)             | Ischaemic 5 (62.5 %)         |
|                                  | Haemorrhagic 3 (37.5 %)      |
| Hemiparetic side, n (%)           | Left 4 (50 %)                |
|                                  | Right 3 (37.5 %)             |
|                                  | Bilateral 1 (12.5 %)         |
| Years post stroke, median (IQR)   | 1 y 42 d (8.2 y)             |
| FAC, mean (range)                 | 1.8 (0–3)                    |
| MMSE score                        | 27.1 ± 3.2                   |
| Comorbidities, n (%)              | Diabetes mellitus 1 (12.5 %) |
|                                  | Hypertension 5 (62.5 %)      |
|                                  | Dyslipidemia 2 (25 %)        |
|                                  | None 3 (37.5 %)              |
| Antihypertensive medications, n (%)| β-blocker 1 (12.5 %)         |
|                                  | ACE inhibitors 3 (37.5 %)    |
|                                  | Calcium channel blockers 1 (12.5 %) |
|                                  | None 3 (37.5 %)              |

Values are mean ± SD unless otherwise indicated
Abbreviations: n, number; SD, standard deviation; MMSE, Mini Mental State Examination; IQR, interquartile range; FAC, Functional Ambulation Category; ACE, angiotensin-converting-enzyme

### Robotics-assisted tilt table (RATT)

A RATT system (Erigo, Hocoma AG, Switzerland) was augmented to facilitate active participation during exercise. The basic RATT is a motorized tilt table with a body harness to support the body and two motor drives to support cyclical movement of the legs. Two thigh cuffs fix the legs and interface to the leg drives, and two foot plates support the feet. The RATT is designed to be used for early rehabilitation in neurological patients to provide early mobilization and intensive sensorimotor stimulation. Additionally, it is also claimed to enhance cardiovascular output by cyclic leg loading. During the therapy, the RATT can be tilted up from 0 to 80° and the cyclic leg movement can be set to a stepping cadence between 8 and 80 steps/min.

With the augmented RATT system, patients are able to see their exercise work rate together with a target work rate [11]. This was achieved by adding individual force sensors to the left and right leg cuffs, a work rate estimation algorithm and a real time visual feedback system (Fig. 1). Patients were instructed to adapt their volitional leg effort to follow the target. Active exercise is achieved by producing forces into the leg cuffs in synchrony with the movement of the RATT.

### Experimental procedures

Patients took part in three exercise sessions, each separated by a minimum of 24 h: a familiarization, an incremental exercise test (IET), and a constant load test (CLT). Patients were instructed to avoid strenuous activity within the 24 h before the test sessions and not to consume a large meal, caffeine or nicotine in the three hours prior to testing [19]. For each test, the patient was first transferred and secured to the RATT in accordance with the manufacturer’s guidelines. Then the additional measurement systems (i.e. automatic blood pressure monitoring, and mask for breath-by-breath gas analysis) were fitted. The patient was then tilted upwards to 60°. During the test, the stepping cadence was set at 80 steps/min.

The familiarization was to instruct patients regarding the RATT, the measurement systems and the test procedures. It included a short ramp phase of 5-min duration and work rate ramp of 3 W/min to allow estimation of an appropriate ramp rate for the subsequent IET.

The IET consisted of: (1) a recorded rest phase, where the patient lay passively on the RATT for 3 min; (2) a passive phase, where the RATT moved the patient’s legs for 5 min; (3) a ramp phase, where the patients actively moved their legs in synchrony with the RATT motion while attempting to follow the linearly increasing work rate target. The work rate ramp was set individually in the range of 1.2 to 3.5 W/min based on observations from the ramp phase of the familiarization with the aim to bring the patient to their functional limit within 8–12 min; and
(4) a recovery phase, similar to the initial passive phase, where the RATT moved the patient’s legs for 5 min. The termination criteria for the ramp phase followed the American College of Sports Medicine guidelines [18]. Additionally, blood pressure (BP) was used as a termination criterion: systolic BP > 210 mmHg or diastolic BP > 115 mmHg [20].

The CLT consisted of: (1) a rest phase for 3 min; (2) a passive phase for 5 min; (3) a constant load phase, where the patient actively moved their legs in synchrony with the RATT motion to follow the constant target work rate (the work rate was set at 40 % of peak work rate ($WR_{peak}$) obtained from the IET) for 10 min; and (4) a recovery phase for 5 min.

**Outcome measures**

Metabolic gas exchange was recorded using a breath-by-breath system (MetaMax 3B, Cortex Biophysik GmbH, Germany) and outcome variables were analysed using the associated Metasoft software. Prior to each test, pressure, volume and gas calibration were performed: volume was calibrated using a 3-L syringe and gas was calibrated using ambient air and a certified precision gas mixture (15 % oxygen and 5 % carbon dioxide) according to the manufacturer’s instructions. Heart rate was continuously recorded using a chest strap (model T34, Polar Electro Oy, Finland). Blood pressure was measured by automatic sphygmomanometry (HEM 907XL, Omron Corporation, USA) every 2 min during the tests.

Outcome measures for IETs were as follows. Peak oxygen uptake ($V'O_2_{peak}$) was determined as the maximum of a 30-s average during the ramp phase. The peak respiratory exchange ratio ($RER_{peak}$) was the 30-s average of RER at the time of $V'O_2_{peak}$. Peak heart rate ($HR_{peak}$) was defined as the highest value of HR reached during the ramp phase. The peak work rate ($WR_{peak}$) was
calculated as the maximum of a 10-s moving average of the recorded work rate.

The first and second ventilatory thresholds (VT and RCP) were determined independently by two experienced raters (JSa and KH) using the methods described by Binder et al. [21] and summarised in the following paragraphs, and the average of the two ratings was taken. The thresholds were characterized by the value of VO\(_2\) at the point where the criteria given below were deemed to be fulfilled. Threshold analysis was carried out visually based on 15-breath averages of the variables concerned.

The first ventilatory threshold, i.e. that which is denoted here as the VT, was determined graphically using the combination of these criteria: (1) the point at which the ventilatory equivalent for oxygen (VE/VO\(_2\)) reaches a minimum or has its first increase without a simultaneous increase in the ventilatory equivalent for carbon dioxide (VE/\(V^{'CO_2}\)); (2) the point at which partial pressure of end-tidal oxygen tension (\(P_{ET}^{'O_2}\)) reaches a minimum or has its first increase without a decrease in the partial pressure of end-tidal carbon dioxide tension (\(P_{ET}^{'CO_2}\)); and, (3) the deflection point of carbon dioxide output (\(V^{'CO_2}\)) versus oxygen uptake (\(V^{'O_2}\); V-slope method). When these 3 criteria gave different results, the first two were prioritized.

The second ventilatory threshold, i.e. the respiratory compensation point (RCP), was determined graphically by inspection of: (1) the minimal value or nonlinear increase of VE/\(V^{'CO_2}\); (2) the turning point of \(P_{ET}^{'O_2}\) and \(P_{ET}^{'CO_2}\); and, (3) the deflection point of minute ventilation (VE) versus \(V^{'CO_2}\) [21]. Again, if these 3 criteria gave different results, the first two were prioritized.

Borg CR10 ratings of perceived exertion (RPE) for dyspnea and leg effort [22] were recorded every 3 min during the tests. The reasons for test termination were recorded.

Outcome measures for CLTs were the steady-state VO\(_2\) and heart rate during the rest, passive and constant load phases. The accuracy of the patient’s achievement of target work rate was quantified by the root mean square error (RMSE) between the target and actual work rates between the first and ninth min of the constant load phase. The Borg CR10 RPE ratings for dyspnea and leg effort were recorded.

Criteria to determine feasibility of the RATT for exercise testing and training

The criteria for feasibility assessment were [23]: (i) implementation (technical feasibility of the augmented RATT for exercise testing), (ii) acceptability (was the exercise tolerable?), and (iii) responsiveness (was there a measurable, high-level cardiopulmonary reaction?). Incremental CPET was considered to have satisfied responsiveness criteria if in a given patient at least one of the following outcomes could be identified from the IETs (adapted from Marzolini et al. [8]): VO\(_{2\text{max}}\), WR\(_{\text{max}}\), VT or RCP.

The first two of these, VO\(_{2\text{max}}\) and WR\(_{\text{max}}\) indicate whether a patient’s functional capacity in terms of cardiopulmonary and/or neuromuscular exertion was reached:

- VO\(_{2\text{max}}\) was deemed to have been achieved if at least one of the following was observed: plateau in VO\(_2\) (increase in VO\(_2\) less than 150 mL in the final minute of exercise [24]), RER \(\geq 1.10\) (or RER \(\geq 1.05\) for age \(\geq 50\) [25]) or HR\(_{\text{peak}}\) \(\geq HR_{\text{max}}\) – 10 [8]. Here, HR\(_{\text{max}}\) was obtained from an age-related prediction formula [18].
- Achievement of WR\(_{\text{max}}\) was marked by a plateau in WR with the patient no longer able to reach the WR target.

Constant-load CPET was considered to have satisfied responsiveness criteria if the intensity and duration of steady-state exercise during CLTs was found to have met current recommendations for exercise and physical activity after stroke; these are defined as 40 to 70 % of VO\(_2\) reserve or HR reserve; or, 55 to 80 % of HR\(_{\text{max}}\); or, RPE of 11 to 14 on the Borg scale (6-20) [4]. VO\(_2\) reserve is defined as VO\(_{2\text{peak}}\) – VO\(_{2\text{rest}}\) and HR reserve as HR\(_{\text{peak}}\) – HR\(_{\text{rest}}\) [18].

Statistical analysis

Descriptive statistics were used to evaluate the distribution of the variables. Continuous variables are presented as mean \(\pm\) standard deviation. Categorical variables are presented as frequencies and percentages. All analyses were performed using SPSS version 19 (IBM Corporation, USA).

Results

(i) Implementation: The augmented RATT could be successfully used to implement both the IET and the CLT in stroke patients without the need to further modify the system. There were no technical problems that interrupted the tests.

(ii) Acceptability: The patients could understand the task to keep up with the work rate target using the visual feedback system and adaptation of their volitional leg effort. All patients could exercise until the end of the protocols without any complications. All tests were completed successfully according to the termination criteria (IET) or pre-specified duration (CLT).

(iii) Responsiveness: IET (\(n = 8\); Table 2): Absolute VO\(_{2\text{peak}}\) was 845 \(\pm\) 266 mL/min (relative VO\(_{2\text{peak}}\) was 11.9 \(\pm\) 4.0 mL/kg/min), which corresponds to
Table 2 Summary of outcome variables from incremental exercise tests (n = 8)

| Outcome variable | Value     | Range      |
|------------------|-----------|------------|
| Peak exercise variables |           |            |
| VO₂peak absolute (mL/min) | 844.8 ± 265.7 | 352.0 – 1176.0 |
| VO₂peak relative (mL/min/kg) | 119 ± 4.0 | 56 – 17.5 |
| VO₂peak as % of predicted VO₂max [14] | 45.2 ± 8.8 | 32.3 – 58.2 |
| HRpeak (beats/min) | 117.3 ± 31.5 | 66.0 – 155.0 |
| VO₂peak as % of predicted HRpeak (220-age) | 72.0 ± 17.7 | 44.9 – 99.3 |
| SBP (mmHg) | 192.9 ± 29.9 | 140.0 – 220.0 |
| DBP (mmHg) | 88.5 ± 12.7 | 78.0 – 110.0 |
| Rate-pressure product | 2300 ± 798 | 924 – 3410 |
| RERpeak | 1.00 ± 0.10 | 0.78 – 1.20 |
| Borg CR10 RPE scale dyspnea | 5.4 ± 2.8 | 1.0 – 9.0 |
| Borg CR10 RPE scale leg effort | 6.6 ± 1.9 | 4.0 – 9.0 |
| Oxygen cost of work (mL/min/W) (n = 6) | 21.9 ± 3.8 | 16.0 – 26.0 |
| WRpeak (W) | 225 ± 13.0 | 55 – 38.6 |
| Time to VO₂peak (sec) | 7425 ± 161.8 | 600.0 – 10800.0 |

Submaximal exercise variables (n = 7)

|                      |            |            |
|----------------------|------------|------------|
| Absolute VO₂ at VT (mL/min) | 677.1 ± 187.1 | 520.0 – 1053.0 |
| Relative VO₂ at VT (mL/kg/min) | 89.2 ± 2.9 | 69 – 15.0 |
| VT as % of VO₂peak | 75.1 ± 17.6 | 523 – 100.8 |
| RER at VT | 0.88 ± 0.08 | 0.78 – 0.96 |
| Absolute VO₂ at RCP (mL/min) (n = 6) | 800.3 ± 206.7 | 547.0 – 1136.0 |
| Relative VO₂ at RCP (mL/kg/min) (n = 6) | 107.0 ± 3.1 | 73 – 16.2 |
| RCP as % of VO₂peak (n = 6) | 84.8 ± 11.6 | 70.6 – 99.3 |
| RER at RCP (n = 6) | 1.02 ± 0.13 | 0.84 – 1.24 |

Values are mean ± SD

Abbreviations: VO₂peak, peak oxygen uptake; VO₂, oxygen uptake; HR, heart rate; HRpeak, peak heart rate; RERpeak, peak respiratory exchange ratio; SBP, systolic blood pressure; DBP, diastolic blood pressure; RPE, rating of perceived exertion; WRpeak, peak work rate; VT, 1st ventilatory threshold; RER, respiratory exchange ratio; RCP, respiratory compensation point.

45.2 % of the expected VO₂max based on the prediction method of Wasserman et al. [14]. HRpeak was 117 ± 32 beats/min, which is 72.0 % of the predicted value (Table 2). WRpeak was 225 ± 13.0 W. The average ratings of perceived exertion (RPE, Borg CR10) for dyspnea and leg effort at the end of the ramp phase were 5.4 and 6.6, respectively; these lie on the qualitative range of “hard” to “very hard”.

Reasons for termination of the IET were: leg fatigue (n = 4, 50 %); abnormal blood pressure, i.e. systolic BP > 210 mmHg, (n = 2, 25 %); breathing effort (n = 1, 12.5 %); and generalized fatigue (n = 1, 12.5 %).

The VT was identified in 7 patients (4 female) and the RCP in 6 patients (3 female). The average absolute VO₂ values at VT and RCP were 677 and 800 mL/min (relative VO₂ at VT and RCP was 8.9 and 10.7 mL/kg/min) which represent 75.1 % (VT) and 84.8 % (RCP) of mean VO₂peak.

Incremental CPET provided sufficient information to satisfy the responsiveness criteria, i.e. VO₂max, WRmax, VT or RCP were able to be determined, in all 8 patients (4 female, 4 male): all 8 patients reached the limit of functional capacity in terms of either VO₂max (7 patients – 4 female) or WRmax (6 patients – 3 female). Of the 7 patients deemed to have satisfied the criteria for VO₂max, 5 reached a plateau, 3 met the RER criterion and 1 met the HRmax criterion.

To illustrate typical IET responses, Fig. 2 shows the target and measured work rates as well as cardiopulmonary responses from the IET in Patient 8. Fig. 3 shows the graphical plots for determination of the VT and the RCP in the same patient.

CLT (n = 8; Table 3): The transition from passive to constant load exercise yielded a higher increase in VO₂ (2.7 mL/kg/min) than did the transition from rest to passive (0.9 mL/kg/min) (Table 3). During the active phase of the exercise, all patients were able to achieve the recommended training intensity level [4] based on percentage of VO₂ reserve, percentage of HRmax and RPE: the constant work rate was set at 40 % of individual WRpeak values which resulted, on average, in a steady-state VO₂ of 49 % of VO₂ reserve, steady-state HR of 56 % of predicted HRmax and RPE > 2. All patients could maintain the active work rate for 10 min as prescribed. The accuracy of maintaining the work rate target (RMSE) was 1.3 W.

Discussion

The aim of this study was to evaluate the feasibility of the augmented RATT for incremental cardiopulmonary exercise testing and exercise training in dependent-ambulatory stroke patients. Feasibility assessment considered technical feasibility, patient tolerability, and cardiopulmonary responsiveness.

Feasibility for incremental cardiopulmonary exercise testing

For all 8 patients tested, incremental CPET provided sufficient information to satisfy the responsiveness criteria, i.e. VO₂max, WRmax, VT or RCP were successfully identified. All 8 patients also reached their limit of functional capacity due to either cardiopulmonary limitations (VO₂max criteria; 7 patients – 4 female) or neuromuscular limitations (WRmax criteria; 6 patients – 3 female). Of these 8 patients, 5 reached both sets of criteria for cardiopulmonary and neuromuscular capacity, 2 patients satisfied only the cardiopulmonary criteria, and 1 patient reached only the neuromuscular limitation.

It is interesting that, in these numbers, female patients are at least as highly represented as males. Marzolini
et al. [8] previously noted that females after stroke were much less likely than males to achieve similar feasibility criteria from baseline CPETs: 40 % for females vs. 81 % for males. That difference was attributed to greater disability and weakness in the females examined in a study of mildly-impaired patients. The results herein, with severely-impaired patients, indicate that such measurement difficulties can be overcome by employing appropriate testing equipment, i.e. the augmented RATT. However, more subjects are required to reliably study these male–female ratios in the outcomes.

The VO\textsubscript{2}peak reported in this study (11.9 ± 4.0 mL/kg/min) is lower than values previously reported in ambulatory stroke patients using: cycle ergometry, 17.2 ± 3.0 mL/kg/min [26]; recumbent cycle ergometry, 16.0 ± 1.2 mL/kg/min [27]; or a treadmill with body weight support, 14.4 ± 5.1 mL/kg/min [24]. This low value may be attributable in part to the more profound disability in the patients in the present study and in part to the observation that, in normal subjects, the RATT VO\textsubscript{2}peak is approximately 20 % lower than with a cycle ergometer and 30 % lower than for a treadmill [28].

HR\textsubscript{peak} was on average 72 % of the predicted value. The rate-pressure product, which reflects the cardiovascular load during exercise, was 230. These results are comparable to previously documented results in mildly to moderately disabled stroke patients [2, 24, 26, 29]. This suggests a similar myocardial work load.

Successful identification of a VT (7 patients – 4 female) and/or an RCP (6 patients – 3 female) from incremental CPET provides an additional means of prescribing exercise intensity. The relative VO\textsubscript{2}peak at the VT (8.9 mL/kg/min) is lower than previous reports [8, 20], which may be due to the same reasons as described above in relation to lower VO\textsubscript{2}peak. The VT as a percentage of VO\textsubscript{2}peak found in this study is in line with other studies which reported values in the range 73.4 to 89.7 % of VO\textsubscript{2}peak [8, 20, 30].

Feasibility for exercise training
Current standards for prescription of exercise intensity for stroke patients are derived from a subset of the main incremental CPET outcomes. For stroke patients, these are [4]: 40 to 70 % of VO\textsubscript{2} reserve or HR reserve; or, 55 to 80 % of HR\textsubscript{max}; or, RPE between 11 and 14 (6 to 20 scale), which corresponds approximately to 2 to 4.5 on the Borg CR10 RPE scale. It is recommended that this intensity level should be reached on 3 to 5 days per week using 20 to 60 min per exercise session or by multiple 10-min sessions.

The CLTs demonstrate that, during the active phase of the exercise, all patients were able to sustain the recommended intensity level based on percentage of VO\textsubscript{2} reserve, percentage of HR\textsubscript{max}, and RPE for 10 min: the constant
work rate was set at 40% of individual WRpeak values which resulted, on average, in a steady-state \( V'O_2 \) of 49% of \( V'O_2 \) reserve, steady-state HR of 56% of HRmax and RPE > 2.

During the passive phases of the CLTs the exercise intensity was far below the recommended levels: passive \( V'O_2 \) was on average 18% of \( V'O_2 \) reserve and passive HR was 9% of HR reserve. This demonstrates that muscle activation is very low during passive movement and emphasises the need for active participation of the patient using the work rate biofeedback screen implemented within the augmented RATT. This low intensity of passive motion confirms a previous report with robotics-assisted treadmill exercise [31]. With this low intensity, patients cannot effectively improve their cardiopulmonary fitness.

These considerations show that the augmented RATT is a feasible platform for implementation of a prescribed exercise training programme where 10-min bouts of exercise form part of the recommendations, but future work is required to investigate the response to a longitudinal training intervention using the RATT.

**Limitations**

The RATT V'O2peak was previously observed to be approximately 20% lower than the cycle ergometer and 30% lower than the treadmill in normal subjects [28], but it is not certain whether these differences would be the same in stroke patients. Further study is needed to address the comparability of the RATT and the standard exercise testing devices (e.g. cycle ergometer) in patients who are capable of using conventional modalities.

It was observed here that the efficiency of work production on the RATT, as characterized by the inverse of the oxygen cost of the work (mean value 21.9 mL/min/W, Table 2), is substantially lower than for cycle or treadmill ergometry. This was also observed previously in able-bodied subjects on the RATT [12]. This is probably due to a combination of factors including the way the muscle groups are activated and the employment of a possibly non-optimal exercise cadence. Here, a cadence of 80 steps/min was used as this is the highest rate allowed by the device employed. To improve efficiency, a higher stepping rate might be desirable, so that lower forces are needed for a given work rate target (because work rate is the product of torque and angular velocity).

The small sample size in this study may limit generalizability of the data describing cardiopulmonary fitness in dependent-ambulatory stroke patients. The V'O2peak in this study may still be an overestimate in relation to the overall population because we did not include patients who had heart disease or who were...
Table 3 Summary of outcome variables from constant load tests (n = 8)

| Outcome variables | Values |
|-------------------|--------|
| Initial rest phase|        |
| VO₂ absolute (mL/min) | 243.9 ± 33.6 |
| VO₂ relative (mL/min/kg) | 3.30 ± 0.43 |
| HR (beats/min) | 73.8 ± 9.8 |
| Passive phase |        |
| VO₂ absolute (mL/min) | 316.6 ± 75.9 |
| VO₂ relative (mL/min/kg) | 4.20 ± 0.78 |
| HR (beats/min) | 76.3 ± 10.0 |
| Constant load phase |        |
| VO₂ absolute (mL/min) | 519.5 ± 117.7 |
| VO₂ relative (mL/min/kg) | 6.9 ± 1.6 |
| VO₂ as % of VO₂ reserve | 48.7 ± 19.1 |
| HR (beats/min) | 90.3 ± 18.6 |
| HR as % of HR reserve | 34.8 ± 32.0 |
| HR as % of HRmax | 56.0 ± 9.9 |
| WR (W) | 11.5 ± 6.1 |
| Borg CR10 RPE scale dyspnea | 2.1 ± 1.0 |
| Borg CR10 RPE scale leg effort | 2.9 ± 0.6 |
| RMSE of WR | 1.3 ± 1.0 |

Values are mean ± SD
Abbreviations: VO₂, oxygen uptake; HR, heart rate; WR, work rate; W, Watts; RPE, rating of perceived exertion; RMSE, root mean square error

not approved by the cardiologist for cardiac safety. Additionally, the mean age of the patients (58.3 years) was slightly lower than the general stroke population reported in high income countries (66 years) [32] or in Switzerland (72.5 years) [33].

Conclusion
The augmented RATT is deemed feasible for incremental cardiopulmonary exercise testing and exercise training in dependent-ambulatory stroke patients: the approach was found to be technically implementable, it was well tolerated by the patients (acceptability), and substantial cardiopulmonary responses were observed (responsiveness).

Abbreviations
CPET: Cardiopulmonary exercise testing; CLT: Constant load test; HR: Heart rate; HRpeak: Peak heart rate; HRmax: Maximal heart rate; IET: Incremental exercise test; PTPCO₂: Partial pressure of end-tidal carbon dioxide tension; PTETCO₂: Partial pressure of end-tidal oxygen tension; RATT: Robotics-assisted tilt table; RCP: Respiratory compensation point; RERpeak: Peak respiratory exchange ratio; RMSE: Root mean square error; RPE: Rating of perceived exertion; VO₂CO₂: Carbon dioxide output; VE: Minute ventilation; VO₂: Oxygen uptake; VO₂max: Maximal oxygen uptake; VO₂peak: Peak oxygen uptake; VT: Ventilatory threshold; WRpeak: Peak work rate.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
All authors made substantial contributions to the study and to manuscript preparation. JSa, KH, TN and CS contributed to the design of the study. JSa, CH, JSc and CS reviewed patients for inclusion and recruited the patients. JSa, CH, JSc and CS carried out the assessments. JSa, CH, JSc and KH performed the data analysis. JSa, CH, JS, CS, TN and KH participated in drafting and critical review of the manuscript. All authors read and approved the final manuscript.

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