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Construct Validation of the Leisure Time Physical Activity Questionnaire for People with SCI  
(LTPAQ-SCI)

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### Abstract

**Study design.** Cross-sectional construct validation study.

**Objectives.** To test the construct validity of the Leisure Time Physical Activity Questionnaire for People with Spinal Cord Injury (LTPAQ-SCI) by examining associations between the scale responses and cardiorespiratory fitness (CRF) in a sample of adults living with spinal cord injury (SCI).

**Setting.** Three university-based laboratories in Canada.

**Methods.** Participants were 39 adults (74% male;  $M$  age:  $42 \pm 11$  years) with SCI who completed the LTPAQ-SCI and a graded exercise test to volitional exhaustion using an arm-crank ergometer. One-tailed Pearson's correlation coefficients were computed to examine the association between the LTPAQ-SCI measures of mild-, moderate-, heavy-intensity and total minutes per week of LTPA and CRF (peak oxygen uptake [ $\dot{V}O_{2\text{peak}}$ ] and peak power output [ $PO_{\text{peak}}$ ]).

**Results.** Minutes per week of mild-, moderate- and heavy-intensity LTPA and total LTPA were all positively correlated with  $\dot{V}O_{2\text{peak}}$ . The correlation between minutes per week of mild intensity LTPA and  $\dot{V}O_{2\text{peak}}$  was small ( $r = .231$ ,  $p = .079$ ) while all other correlations were medium-sized ( $r$ s ranged from .276 to .443,  $ps < .05$ ). Correlations between the LTPAQ-SCI variables and  $PO_{\text{peak}}$  were also positive but small ( $r$ s ranged from .087 to .193,  $ps > .05$ ), except for a medium-sized correlation between heavy-intensity LTPA and  $PO_{\text{peak}}$  ( $r = .294$ ,  $p = .035$ ).

**Conclusions.** People with SCI who report higher levels of LTPA on the LTPAQ-SCI also demonstrate greater levels of CRF. These results provide further support for the construct validity of the LTPAQ-SCI as a measure of LTPA among people with SCI.



24 and validity<sup>7,8,12</sup>. Using a structured, standardized interview format, respondents are cued to  
25 recall and rate the intensity of all LTPA and activities of daily living (ADL) that they have  
26 performed over the previous 3 days<sup>13</sup>. The PARA-SCI has demonstrated positive evidence of  
27 criterion validity (using both indirect calorimetry and doubly-labeled water as criteria), construct  
28 validity and test-retest reliability<sup>11,14,12</sup>. However, because the PARA-SCI was designed to  
29 capture the types, frequencies, intensities and durations of *all* physical activities, it can create  
30 unnecessary participant and clinician/researcher burden in situations where investigators are  
31 interested only in measuring LTPA<sup>7</sup>. In response to these concerns, the Leisure Time Physical  
32 Activity Questionnaire for People with SCI (LTPAQ-SCI) was developed<sup>15</sup>.

33         The LTPAQ-SCI is an SCI-specific, self-report assessment of LTPA that measures the  
34 number of minutes of mild, moderate, and heavy intensity LTPA that a person performed over  
35 the previous 7 days<sup>15</sup>. It can be self- or interviewer-administered in less than 5 minutes. The  
36 reporting format used in the LTPAQ-SCI parallels the reporting structure of one of the most  
37 widely used self-report measures of PA in the general population--the International Physical  
38 Activity Questionnaire-Short Form<sup>16</sup>.

39         Research has produced positive evidence of the LTPAQ-SCI's test-retest reliability.  
40 Intraclass correlation coefficients were significant for LTPAQ-SCI measures of mild, moderate,  
41 heavy and total LTPA over a one-week test-retest period<sup>15</sup>. A recent study of the test-retest  
42 reliability of a Canadian-French version of the questionnaire produced similarly strong ICCs<sup>17</sup>.  
43 Evidence of the measure's criterion validity was shown by significant correlations between  
44 LTPAQ-SCI measures of mild, moderate, heavy and total LTPA minutes per week and PARA-  
45 SCI measures (i.e., the criterion) LTPA minutes per day at these same intensities<sup>15</sup>.

46 Support for the LTPAQ-SCI's construct validity has been generated in hypothesis-testing  
47 studies<sup>18</sup>. For example, LTPAQ-SCI measures of LTPA have been shown to increase  
48 significantly in response to LTPA-enhancing interventions delivered to adults with SCI<sup>19</sup> and  
49 multiple sclerosis<sup>20</sup>. LTPAQ-SCI measures of LTPA have also been shown to differ in predicted  
50 directions between adults with SCI with low versus high depressive symptomatology,<sup>21</sup> and  
51 between athletes with disabilities who participate in sport at lower (recreational, developmental)  
52 versus higher (provincial, state, national) competitive levels<sup>22</sup>. It is important to note, however,  
53 that construct validation is an ongoing process, and no one single experiment can 'prove'  
54 construct validity<sup>18</sup>. Rather, each supportive study serves to strengthen the construct's  
55 nomological network,<sup>23</sup> by demonstrating that the construct operates predictably within a system  
56 of key concepts.

57 Cardiorespiratory fitness (CRF) is a key concept in relation to LTPA. It is well-established  
58 that participation in moderate- to heavy-intensity exercise (a specific type of LTPA) imparts  
59 significant improvements in the CRF of adults with SCI<sup>1,24</sup>. If the LTPAQ-SCI is to be used as a  
60 measure of LTPA, then its construct validation should include tests of its associations with CRF  
61 (these types of tests are sometimes referred to as tests of 'convergent validity'<sup>18</sup>). Therefore, the  
62 purpose of the present study was to examine the association between the number of minutes per  
63 week of mild, moderate, heavy and total LTPA reported by adults with SCI who completed the  
64 LTPAQ-SCI, and their CRF. It was hypothesized that number of minutes per week of LTPA  
65 would be positively correlated with participants' CRF.

## 66 **Method**

### 67 **Participants**

68 Participants were 51 individuals who completed the LTPAQ-SCI and CRF assessment during  
69 baseline testing for CHOICES (NCT01718977), a multicentre, randomized controlled clinical  
70 trial assessing the effects of two different exercise interventions on cardiovascular health  
71 outcomes in adults with SCI<sup>25</sup>. This construct validation study was planned *a priori*, as a sub-  
72 study within CHOICES, when the trial protocol was designed. CHOICES study inclusion criteria  
73 were: male or female; 18-60 years of age; chronic (>1 year since injury), traumatic, motor-  
74 complete SCI [American Spinal Injury Association Impairment Scale (AIS) A and B]; and  
75 neurological level of injuries (NLI) between the cervical fourth and thoracic sixth vertebrae (C4-  
76 T6). AIS and NLI were determined using the International Standards for neurological  
77 Classification of SCI<sup>26</sup>. Participants were excluded if they had: any medical history of symptoms  
78 of cardiovascular disease; major trauma or surgery in the last six months; fracture within the  
79 previous 12 months; or any psychological or cognitive dysfunction that prevented understanding  
80 English instructions. All study procedures were approved by the research ethics board at each  
81 trial site and all participants provided written informed consent prior to any of the study  
82 procedures.

### 83 **Measures**

84 *LTPAQ-SCI*. The LTPAQ-SCI was administered during an interview conducted by a  
85 research assistant (face-to-face interview at two sites and telephone interview at one site).  
86 Consistent with the LTPAQ-SCI administration instructions,<sup>15</sup> participants were first presented  
87 with a standardized definition of LTPA: “physical activity that you choose to do during your free  
88 time, such as exercising, playing sports, gardening, and taking the dog for a walk (necessary  
89 physical activities such as physiotherapy, grocery shopping, pushing/wheeling for transportation  
90 are not considered LTPA).” Next, participants were given a validated,<sup>11</sup> SCI-specific definition

91 of mild-intensity LTPA and were asked to recall a) the number of days, over the past 7 days, that  
92 they did mild-intensity LTPA and b) on those days, how many minutes they usually spent doing  
93 mild-intensity LTPA. These steps were repeated for moderate-intensity and heavy-intensity  
94 LTPA. The number of minutes per week of LTPA performed at each intensity (mild, moderate  
95 and heavy) was calculated by multiplying the days of activity by the minutes of activity. Total  
96 LTPA was calculated as a sum of LTPA at each intensity, thus yielding the total number of  
97 minutes of LTPA undertaken in the past week.

98 *Cardiorespiratory fitness (CRF)*. All participants underwent an incremental exercise test  
99 using an electronically braked arm-crank ergometer (Lode BV, Groningen, The Netherlands;  
100 Vancouver site, Monark 881E, Monark Exercise AB, Vansbro, Sweden; Toronto and Hamilton  
101 sites) until the point of volitional exhaustion. Heart rate was recorded continuously using a chest  
102 strap HR monitor (T31; Polar Electro Inc., Woodbury, NY, USA). Respiratory gases were  
103 collected using a metabolic cart that was calibrated, prior to each use, according to the  
104 manufacturer's instructions (Parvomedics Truemax 2400, Sandy, Utah, USA; Vancouver site:  
105 Vmax Encore, SensorMedics, California, USA; Toronto site: Moxus Metabolic System, AEI  
106 Technologies, Illinois, USA; Hamilton site).

107 Participants were asked to empty their bladders prior to the test to minimize the influence  
108 of autonomic dysreflexia. The test protocol began with a warm-up of arm cranking at 0 Watts for  
109 two minutes. Afterwards, the protocol continued with 1-minute stages, with a resistance  
110 increment of 5-10 Watts per stage depending on the participant's neurological level of injury<sup>27</sup>.  
111 Participants were instructed to maintain a cycling cadence of 50 revolutions per minute (rpm)  
112 throughout the duration of the test with continuous motivation delivered by the assessor. The test  
113 continued to the point of volitional exhaustion or when the cadence dropped below 30 rpm.



114 Borg's rating of perceived exertion (RPE) 6-20 was administered at the end of every stage<sup>28</sup>.  
115 The highest  $\dot{V}O_2$  of 20-second averaging during the test was recorded as peak oxygen uptake  
116 ( $\dot{V}O_{2\text{peak}}$ ). The highest power output maintained for at least 20 seconds was recorded as peak  
117 power output ( $PO_{\text{peak}}$ ).

## 118 **Procedure**

119 At two sites (Hamilton and Vancouver), the LTPAQ-SCI was administered during the  
120 baseline testing session, prior to the CRF test. At one site (Toronto), the LTPAQ-SCI was  
121 administered 8 days after the fitness test but before starting exercise in the CHOICES trial. This  
122 timing was deliberate to avoid participants reporting any LTPA that was performed as part of the  
123 CHOICES baseline testing or training protocols.

## 124 *Data Management and Analyses*

125 The respiratory exchange ratio (RER) was used to corroborate attainment of  $\dot{V}O_{2\text{peak}}$   
126 during the fitness test. Analyses were conducted only on participants who exhibited an RER  $\geq$   
127 1.00. People with tetraplegia cannot achieve the same  $\dot{V}O_{2\text{peak}}$  and  $PO_{\text{peak}}$  as people with  
128 paraplegia due to more severe autonomic and upper-body motor impairments<sup>29</sup>. Consequently,  
129 the distributions of these values differ for people with tetraplegia versus paraplegia<sup>29</sup>. Therefore,  
130 the measures of CRF (i.e.  $\dot{V}O_{2\text{peak}}$  and  $PO_{\text{peak}}$ ) were standardized for lesion level (i.e., paraplegia  
131 or tetraplegia) through transformations to z-scores prior to analysis.

132 Descriptive statistics were calculated as means, standard deviations, medians and  
133 minimum-maximum for continuous variables, and as percentages for the categorical variables.  
134 Shapiro Wilk tests were used to check the normality assumption. Because the LTPAQ-SCI  
135 variables presented significant deviations from the normal distribution, a square root  
136 transformation was carried out on these variables. Using the transformed variables, one-tailed

137 Pearson correlation coefficients were calculated between the LTPAQ-SCI measures of mild,  
138 moderate, heavy and total LTPA and the measures of CRF (i.e.  $\dot{V}O_{2\text{peak}}$  and  $PO_{\text{peak}}$ ). One-tailed  
139 tests were used given the directionality of the hypotheses. All analyses were conducted using  
140 IBM SPSS Statistics v. 26. Alpha was set at .05 for all analyses given the *a priori* nature of the  
141 hypotheses. Cohen's conventions were used to interpret the magnitude of the correlations (i.e., *r*s  
142 of .10, .30, .50 constitute small, medium and large correlations, respectively)<sup>30</sup>.

## 143 **Results**

### 144 **Preliminary analyses**

145 After excluding data from 12 participants who did not achieve  $RER \geq 1.00$ , 39  
146 participants remained for the main analyses. Excluded participants presented significantly lower  
147  $PO_{\text{peak}}$  and  $VO_{2\text{peak}}$  values than the included ones. In addition, all excluded participants had  
148 tetraplegia. No significant differences were found between excluded and included participants  
149 regarding sex, age, age at injury, time since injury, body mass or height, or LTPAQ-SCI values.  
150 Subsequent analyses were conducted with data from the remaining 39 participants. Table 1  
151 shows the demographic data for both the full sample and the final sample, as well as the *p*-values  
152 of the tests performed to detect potential differences between included and excluded participants

### 153 **Correlations between the LTPAQ-SCI measures of LTPA and aerobic fitness**

154 Table 2 presents the full correlation matrix. Minutes per week of mild-, moderate- and  
155 heavy-intensity LTPA and total LTPA were all positively correlated with  $\dot{V}O_{2\text{peak}}$ . The  
156 correlation between minutes per week of mild intensity LTPA and  $\dot{V}O_{2\text{peak}}$  was small ( $r = .231$ ,  $p$   
157  $= .079$ ) while all other correlations were medium-sized (*r*s ranged from .276 to .443,  $ps < .05$ ).

158 Correlations between the LTPAQ-SCI variables and  $PO_{peak}$  were also positive. However, the  
159 correlations were generally trivial to small ( $r$ s ranged from .087 to .193,  $p$ s > .05) except for the  
160 correlation between heavy-intensity LTPA and  $PO_{peak}$  ( $r = .294, p = .035$ ).

## 161 **Discussion**

162 The purpose of this study was to conduct a test of the construct validity of the LTPAQ-SCI.  
163 As hypothesized, minutes per week of LTPA reported on the LTPAQ-SCI were positively  
164 correlated with participants' CRF. Correlations tended to be stronger for heavy versus mild-  
165 intensity LTPA and for  $\dot{V}O_{2peak}$  than for  $PO_{peak}$ .

166 Overall, the pattern and size of the correlations were similar to correlations reported between  
167 CRF and other self-report measures of PA for people with and without SCI. For instance, in tests  
168 of the PARA-SCI's construct validity,<sup>14</sup> correlations between CRF and moderate- and heavy-  
169 intensity LTPA were medium-sized, while the correlation between CRF and mild-intensity  
170 LTPA was small. These findings align with research demonstrating that in order to produce  
171 significant CRF benefits, adults with SCI must exercise at a moderate- to heavy-intensity<sup>1</sup>.  
172 Exercise of a mild intensity is insufficient<sup>31</sup>. Our results show that the LTPAQ-SCI does indeed  
173 capture CRF-enhancing LTPA in adults with SCI.

174 Our results are also similar to the medium-sized correlations reported in validation studies of  
175 the IPAQ-SF, one of the most widely-used self-report measures of PA for the general population.  
176 For instance, across three studies that reported correlations between the IPAQ-SF measure of  
177 total minutes per week of PA and  $\dot{V}O_{2max}$ , the median correlation was  $r = .30$ <sup>32</sup>. We found a  
178 correlation of  $r = .33$  between LTPAQ-SCI total LTPA and  $\dot{V}O_{2peak}$ . It is worth noting that only  
179 ~50% of the variance in CRF can be explained by environmental factors, such as physical  
180 activity, with the rest attributed to hereditary/genetic factors<sup>33</sup>. Furthermore, additional variance

181 in CRF within the SCI population can be attributed to the severity and exact level of neurological  
182 injury sustained, contributing to the degree of autonomic and functional impairment<sup>34</sup>. Thus, it is  
183 encouraging to observe similar, if not slightly better, associations between LTPA and  $\dot{V}O_{2peak}$  in  
184 individuals with high-level SCI, supporting the construct validity of the LTPAQ-SCI in the  
185 context of other well-used self-report measures of PA.

186 Our analyses suggested that LTPA was more strongly correlated with  $\dot{V}O_{2peak}$  than  $PO_{peak}$ .  
187 This finding differs from results from the PARA-SCI validation studies in which CRF tended to  
188 be more strongly correlated with  $PO_{peak}$  than  $\dot{V}O_{2peak}$ <sup>14</sup>. These discrepancies are likely a statistical  
189 artefact. There was greater variability in  $PO_{peak}$  values in the PARA-SCI validation study than in  
190 the present study. When data variability is reduced, correlations may be lower than expected<sup>35</sup>.  
191 Nevertheless, as the correlations with  $PO_{peak}$  were all positive, and stronger for moderate- and  
192 heavy-intensity LTPA than mild-intensity LTPA, we take this as further support for the construct  
193 validation of the LTPAQ-SCI as a measure of CRF-enhancing LTPA.

194 Importantly, scale validation studies do not confirm that the scale itself is valid. No study can  
195 ‘validate’ a scale. Rather, validation studies substantiate the inferences that can be made about  
196 people based on their scale scores (e.g., the amount of LTPA they do each week)<sup>18</sup> and that the  
197 scale is valid for use with a particular group of people in a particular context<sup>18</sup>. The present study  
198 was conducted with a sample of men and women with chronic, motor complete cervical or high  
199 thoracic injuries. Although we would expect the results to generalize to individuals with  
200 incomplete, lower-level injuries,<sup>14</sup> this hypothesis should be tested in heterogenous samples.

201 There is also a need to conduct LTPAQ-SCI validation studies in countries other than  
202 Canada, because definitions of LTPA may differ across cultural contexts<sup>36</sup>. For instance, the  
203 instructions for completing the LTPAQ-SCI stipulate that physiotherapy should not be counted.

204 This stipulation is included because during development of the PARA-SCI and LTPAQ-SCI,  
205 many of the physiotherapy activities reported by Canadians with SCI, were neither leisure-time  
206 nor fitness-enhancing activities (e.g., passive stretching, practicing transfers, practicing using  
207 mobility equipment)<sup>13</sup>. However, in other countries or contexts, physiotherapy may routinely  
208 include exercise or sport activities and may therefore be counted as LTPA. In a similar vein,  
209 active transportation is uncommon among Canadians with SCI<sup>37</sup> because climate, terrain and  
210 long distances are significant barriers. In some countries, however, it may be more common for  
211 people with SCI to use active forms of transportation (e.g., handcycling in European countries<sup>38</sup>)  
212 in order to get exercise. In these circumstances, it may make sense to report such activities on the  
213 LTPAQ-SCI. By testing the relationships between CRF and LTPAQ-SCI scores, including and  
214 excluding physiotherapy and active transportation activities, users of the LTPAQ-SCI can better  
215 define and measure LTPA in their contexts.

216 Strengths of this study include standardized administrations of the LTPAQ-SCI and the  
217 CRF test, as well as multi-site data collection to maximize participant enrolment. A limitation is  
218 that only one aspect of physical fitness was measured. Muscular strength and endurance are two  
219 additional physical fitness aspects that should correlate positively with LTPAQ-SCI scores<sup>14</sup> and  
220 should be examined in future construct validation studies. Furthermore, if study participants  
221 engaged primarily in strength-training LTPA (e.g., lifting weights), the correlation between their  
222 LTPAQ-SCI measure of minutes per week of LTPA and their CRF may have been attenuated  
223 relative to individuals who engaged primarily in CRF-enhancing LTPA (e.g., arm cycling).

224 Another study limitation is that data collected from nearly half of the tetraplegic  
225 participants (12 out of 27) could not be used because they terminated the CRF test before  
226 achieving criteria indicative of a peak exercise test (i.e.,  $RER \geq 1.00$ ). Because of arm fatigue

227 during exercise testing, peripheral ratings of perceived exertion increase much faster in those  
228 with tetraplegia than paraplegia<sup>39</sup> prompting participants to terminate the test before achieving  
229 peak. Given this challenge, researchers should consider other feasible, valid measures of CRF  
230 that could be used in LTPAQ-SCI construct validation studies involving participants with  
231 tetraplegia. An alternative construct validation approach may be to assess associations between  
232 LTPAQ-SCI scores and 7-day overall physical activity levels measured via wearable devices.  
233 While limitations of accelerometers attached to a single anatomical location or wheelchair have  
234 been noted in people with SCI<sup>7,9</sup>, the estimation of physical activity intensity can be improved by  
235 utilising multi-sensor devices that incorporate physiological signals (such as galvanic skin  
236 responses or heart rate) and utilising complex or individualised modelling approaches<sup>40,41</sup>.  
237 Combined with the use of diaries or logs to distinguish periods of LTPA from other physical  
238 activity types, assessing the associations between outputs from multi-sensor wearable devices  
239 and the LTPAQ-SCI may be a way to test the validity of this measure while overcoming some of  
240 the challenges noted with assessing CRF in individuals with tetraplegia.

241 In conclusion, the results of the present study demonstrate that self-reported LTPA, as  
242 measured by the LTPAQ-SCI, is positively correlated with CRF in adults with chronic, motor  
243 complete cervical or high thoracic SCI. When considered with previous research showing that  
244 LTPAQ-SCI scores vary in predictable ways across meaningful groups and in response to  
245 behavioural interventions<sup>15,17-22</sup>, these results provide further support for the construct validity of  
246 the LTPAQ-SCI as a measure of LTPA for adults with SCI. Further construct validation studies  
247 are needed to demonstrate the validity of the LTPAQ-SCI for use as a measure of LTPA in more  
248 heterogeneous samples of people with SCI and in other countries and contexts.

249           **Data archiving:** The datasets generated during and/or analysed during the current study  
250 are available from the corresponding author on reasonable request.

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255 Health Network (12-5797).

256           **Conflict of interest:** The authors declare that they have no conflict of interest.

257           **Author contributions:** KAMG was responsible for conceptualizing and designing the  
258 study, interpreting results and writing the report. JU-C was responsible for analyzing the data,  
259 writing the results, creating tables and providing feedback on the report. AAA was responsible  
260 for collecting and cleaning the data, drafting the methods section and providing feedback on the  
261 report. TEN was responsible for assisting during data collection, drafting the methods section,  
262 assisting with data interpretation, and providing feedback on the report. JSA was responsible for  
263 assisting in the design of the study protocol, drafting the methods section, assisting with data  
264 interpretation, and providing feedback on the report. KDC was responsible for assisting during  
265 data collection and providing feedback on the report. MH was responsible for assisting during  
266 data collection and providing feedback on the report. AK is the Principal Investigator for the  
267 CHOICES study and was responsible for designing and overseeing implementation of all aspects  
268 of the CHOICES protocol and providing feedback on the report.

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Table 1. Characteristics of Participants in the Full Sample and the Analyzed Sample and p-values of Tests to Detect Differences Between Included and Excluded Participants.

|                              | Full Sample (n=51)                          | Analyzed Sample (n=39)                      | p-value |
|------------------------------|---|---|---------|
|                              | n (% total); M $\pm$ SD;<br>Median, min-max | n (% total); M $\pm$ SD;<br>Median, min-max |         |
| Sex                          |   |   | .964    |
| Male                         | 38 (74%)                                    | 29 (74%)                                    |         |
| Female                       | 13 (26%)                                    | 10 (26%)                                    |         |
| Age (years)                  | 42 $\pm$ 10<br>43, 22-60                    | 42 $\pm$ 10<br>43, 22-60                    | .638    |
| Age at injury (years)        | 28 $\pm$ 13<br>24, 3-57                     | 29 $\pm$ 14<br>25, 3-57                     | .252    |
| Years post-injury            | 14 $\pm$ 11<br>10, 1-42                     | 13 $\pm$ 11<br>10, 1-42                     | .502    |
| Level and severity of injury |   |   | <0.001  |
| Tetraplegia AIS A            | 16 (31%)                                    | 8 (21%)                                     |         |
| Tetraplegia AIS B            | 11 (22%)                                    | 7 (18%)                                     |         |
| Paraplegia AIS A             | 24 (47%)                                    | 24 (61%)                                    |         |
| Body mass (kg)               | 78.5 $\pm$ 17.6<br>78.2, 44.9-135.7         | 79.2 $\pm$ 17.3<br>78.5, 44.9-135.7         | .764    |
| Height (cm)                  | 176 $\pm$ 10<br>177, 158-200                | 175 $\pm$ 8<br>176, 158-188                 | .201    |
| VO <sub>2peak</sub>          | 12.53 $\pm$ 5.48<br>11.24, 5.49-29.84       | 13.93 $\pm$ 5.49<br>13.60, 6.07-29.84       | <.001   |
| PO <sub>peak</sub>           | 52 $\pm$ 29<br>50, 10-130                   | 60 $\pm$ 28<br>60, 10-130                   | .001    |
| Mild LTPA                    | 204 $\pm$ 278<br>120, 0-1680                | 221 $\pm$ 308<br>135, 0-1680                | .555    |
| Moderate LTPA                | 102 $\pm$ 118<br>60, 0-480                  | 115 $\pm$ 124<br>60, 0-480                  | .093    |
| Heavy LTPA                   | 56 $\pm$ 100<br>15, 0-480                   | 60 $\pm$ 96<br>20, 0-480                    | .260    |
| Total LTPA                   | 363 $\pm$ 395<br>240, 0-2405                | 395 $\pm$ 431<br>240, 0-2405                | .291    |

AIS: ASIA Impairment Scale

Table 2. Correlation Matrix Showing Pearson Correlation Coefficients for Cardiorespiratory Fitness and LTPAQ-SCI Measures of Mild, Moderate, Heavy and Total Leisure-Time Physical Activity (LTPA)

| <i>Measure</i>                  | VO <sub>2peak</sub> | PO <sub>peak</sub> | Mild LTPA | Moderate LTPA | Heavy LTPA | Total LTPA |
|---------------------------------|---------------------|--------------------|-----------|---------------|------------|------------|
| VO <sub>2peak</sub> (mL/kg/min) | 1                   |                    |           |               |            |            |
| PO <sub>peak</sub> (Watts)      | .773**              | 1                  |           |               |            |            |
| Mild LTPA (min/wk)              | .231                | .087               | 1         |               |            |            |
| Moderate LTPA (min/wk)          | .276*               | .193               | .315*     | 1             |            |            |
| Heavy LTPA (min/wk)             | .443**              | .294*              | .225      | .499**        | 1          |            |
| Total LTPA (min/wk)             | .330*               | .176               | .815**    | .729**        | .591**     | 1          |

Note. \*p<0.05; \*\*p<0.01 (one-tailed).

LTPAQ-SCI is the Leisure Time Physical Activity Questionnaire-Spinal Cord Injury. VO<sub>2peak</sub> is peak volume of oxygen consumption and PO<sub>peak</sub> is peak power output during the cardiorespiratory fitness test.