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Individual calibration of accelerometers in children and their health-related implications

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1 <u>Title Page</u>

- 2 <u>Title:</u> Individual calibration of accelerometers in children and their health-related
- 3 implications
- 4 **<u>Running Title:</u>** Individual PA calibration and health implications
- 5 **Keywords:** physical activity, accelerometry, threshold, children

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- 58

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61 62

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72 Abstract

73 This study compared children's physical activity (PA) levels, the prevalence of children meeting current guidelines of ≥60 minutes of daily moderate to vigorous PA (MVPA), and 74 75 PA-health associations using individually calibrated (IC) and empirical accelerometer 76 cutpoints. Data from 75 (n = 32 boys) 10-12 year old children were included in this study. Clustered cardiometabolic (CM) risk, directly measured cardiorespiratory fitness (CRF), 77 anthropometric and 7 day accelerometer data were included within analysis. PA data were 78 79 classified using Froude anchored IC, Evenson et al., 2008 (Ev) and Mackintosh et al., 2012 80 (Mack) cutpoints. The proportion of the cohort meeting ≥60mins MVPA/day ranged from 81 37%-56% depending on the cutpoints used. Reported PA differed significantly across the cutpoint sets. IC LPA and MPA were predictors of CRF (LPA: standardised $\beta = 0.32$, p = 82 0.002, MPA: standardised $\beta = 0.27$ p = 0.013). IC MPA also predicted BMI Z-score 83 84 (standardised β = -0.35, p = 0.004). Ev VPA was a predictor of BMI Z-score (standardised β = -0.33, p = 0.012). Cutpoint choice has a substantial impact on reported PA levels though no 85 significant associations with CM risk were observed. Froude IC cut points represent a 86 promising approach towards classifying children's PA data. 87

88 Introduction

Regular participation in physical activity (PA) in childhood is associated with reduced 89 cardiometabolic risk (Andersen, Riddoch, Kriemler, & Hills, 2011), improved bone health 90 (Boreham & McKay, 2011), reduced adiposity (McMurray & Ondrak, 2013), and improved 91 92 psychological well-being (Biddle & Asare, 2011). PA guidelines state that children should 93 accrue at least 60 minutes of daily moderate to vigorous PA (MVPA) to receive health benefits 94 (WHO, 2010). The accurate measurement of PA is essential to investigate the associations 95 between PA and health, estimate the prevalence of inactivity, and identify children in need of intervention. Accelerometry is the most commonly used objective method for assessing free-96 97 living PA in children, and has acceptable validity and reliability (Cain, Sallis, Conway, Van 98 Dyck, & Calhoon, 2013). Despite this, no consensus exists with regards to the treatment of 99 accelerometer data and inconsistent use of cutpoints presents challenges when quantifying the 100 prevalence of inactivity (Ekelund, Tomkinson, & Armstrong, 2011), making comparisons 101 between studies (Hislop, Bulley, Mercer, & Reilly, 2012) and establishing the relationship 102 between PA and health outcomes (Bailey, Boddy, Savory, Denton, & Kerr, 2013).

103

104 Previous research has compared the classification accuracy of published thresholds (PTs) in 105 youth using calibration studies, and recommended that researchers use Evenson's (Evenson, 106 Catellier, Gill, Ondrak, & McMurray, 2008) cutpoints to classify children's PA (Trost, 107 Loprinzi, Moore, & Pfeiffer, 2011). However, the empirical cutpoints examined by Trost et 108 al. (2011) applied universal cutpoints to all children, with only one age-specific cutpoint 109 included in the analysis (Freedson, Pober, & Janz, 2005). Such cutpoints fail to account for wide variations in accelerometer counts observed between children when engaging in PA at 110 equivalent intensities (Rowlands, 2007). Subsequently researchers have proposed the use of 111 individually calibrated (IC) approaches to improve the classification of children's PA 112 (Mackintosh, Fairclough, Stratton, & Ridgers, 2012). 113

One method of deriving individual cutpoints, particularly when using hip-mounted 115 accelerometers, is to adjust cutpoints to account for limb length using relevant biomechanical 116 117 theory. The Froude (Fr) number (Minetti, 2001) offers one solution to the standardisation of 118 cutpoints for individuals, by taking the length of a given characteristic, in this case leg length, into account. Froude numbers are calculated using the equation: $Fr = v^2/g^{*1}$ where 'v' is the 119 speed, 'g' represents gravitational acceleration and 'l' is the length of the characteristic. The 120 theory of dynamic similarity suggests that geometrical bodies have similar gait dynamics if 121 122 the Fr number is kept constant (Alexander, 1989). For example, the Fr number of 0.25 represents optimum walking speed and Fr 0.5 is the point at which running occurs in most 123 bipedal bodies, including humans (Kram, Domingo, & Ferris, 1997). Therefore for a given Fr 124 125 number and related walking speed gait dynamics should be relatively consistent between 126 participants, which in turn allows for a simple method of creating individualised and 127 comparable thresholds. Despite its potential utility, few studies have utilised the Froude 128 approach to individually calibrate accelerometer cutpoints (Boddy et al., 2014). As PA is 129 positively associated with health, the potential to better examine these relationships with more 130 precise estimates of PA is important, especially as relationships between PA and variables 131 such as CRF and adiposity are often weaker than may be expected. To date no have examined 132 differences in reported PA or PA-health associations between IC and empirically derived 133 group level cutpoints. Therefore, the aims of this study were to compare children's physical 134 activity (PA) levels, the prevalence of children meeting current guidelines of 60 minutes of 135 daily moderate to vigorous PA (MVPA), and PA-health associations using individually calibrated Fr (IC) and empirical accelerometer cutpoints. 136

137

138 <u>Methods</u>

139 Participants and Settings

The data for this analysis were taken from the REACH Year 6 study (Boddy et al., 2014).
Seventy-five children (n = 32 boys) 10-12 years of age agreed to take part in the study which
had ethical clearance from the respective institutional ethics committees. The study was

143 conducted in Liverpool, England (2010, n = 39) and Belfast, Northern Ireland (2011, n = 35).

144 Each participant attended one school-based blood sampling session and one laboratory testing

- session. Participants also wore an accelerometer to quantify PA over seven days.
- 146
- 147
- 148 Procedure and Measurements

149 Anthropometrics: Stature, sitting stature to the nearest 0.1cm (Seca Ltd. Birmingham, UK) 150 and body mass to the nearest 0.1kg (Seca Ltd. Birmingham, UK) were assessed using standard 151 techniques (Lohman, Roche, & Martorell, 1988). Waist circumference was measured to the nearest 0.1cm. Body mass index (BMI), BMI Z-scores (Cole, Freeman, & Preece, 1995) and 152 153 somatic maturation (Mirwald, Baxter-Jones, Bailey, & Beunen, 2002) were calculated. High-154 resolution ultrasound (Terason, t3000; Aloka, London, UK) was used to assess flow mediated 155 dilation (FMD) and % FMD calculated using the equation: ((Peak artery diameter - Baseline 156 artery diameter)/ Baseline artery diameter)*100) was calculated. Blood pressure (BP) was 157 measured on the left arm after 15mins rest in a supine position using an automated BP monitor 158 (Omron Healthcare UK Limited, Miton Keynes, UK).

159

160 Cardiorespiratory fitness assessment: After treadmill familiarisation, participants completed 161 an individually calibrated continuous, incremental (2mins stages) treadmill (both sites: HP 162 Cosmos, Traunstein, Germany) protocol to volitional exhaustion using online gas analysis 163 (Liverpool: Jaeger Oxycon Pro, Viasys Health Care, UK, Ulster: COSMED, Quark, Italy) to 164 measure peak oxygen uptake (VO_2 peak). Treadmill speeds for the first two stages of the test 165 were anchored to Froude numbers 0.25 (MPA) and 0.5 (VPA) for each participant. For this study leg length was used as the characteristic. An example equation to calculate treadmill 166 speed for an individual with a leg length of 0.67m for a Fr number of 0.25 would be: treadmill 167 speed (m/s) = $\sqrt{(0.25^{*}(9.81^{*}\ 0.67))}$, which would result in a speed of 1.28 m/s or 4.61 km/h. 168 Participants wore an ActiGraph accelerometer (ActiGraph GT1M, MTI Health Services, 169 170 Pensacola, FL) at the right hip and heart rate monitor (Polar Electro Oy, Kempele, Finland)

set to record using 5 second epochs throughout the treadmill protocol. The highest 15-second

- average oxygen uptake was used to represent VO₂peak (ml/kg/min) for each participant.
- 173

174 Blood sampling: On a different day to the laboratory visits, children attended their school sites 175 to provide a fasting venous blood sample. Experienced phlebotomists obtained ~ 10ml of 176 venous blood following an overnight fast. Samples were taken between 8.30 and 10.30am. 177 After providing a sample children were given breakfast. Blood samples were transported to 178 the pathology laboratories at Alder Hey Children's Foundation NHS Trust or the Ulster 179 Hospital for analysis. Blood was analysed for triglycerides, cholesterol, high density 180 lipoprotein cholesterol (HDL-c), glucose, adiponectin, and high sensitivity C-reactive protein 181 (CRP) using assay methods that were standardised between sites. Blood markers were used in 182 combination with FMD%, blood pressure and waist circumference to calculate a clustered 183 cardiometabolic risk score by standardising individual risk components and summing them to 184 create a continuous clustered risk variable. This approach has been used in several similar 185 studies (Andersen, Hasselstrom, Gronfeldt, Hansen, & Karsten, 2004; Anderssen et al., 2007; 186 Boddy et al., 2014; Buchan, Young, Boddy, & Baker, 2014).

187

188 Physical activity assessment: Children wore an ActiGraph (ActiGraph GT1M) uniaxial 189 accelerometer on their right hip during waking hours for seven consecutive days. The monitors 190 recorded activity using 5 second epochs to account for the sporadic nature of children's 191 physical activity (Baquet, Stratton, Van Praagh, & Berthoin, 2007). Periods of 20 minutes of 192 consecutive zero counts (1 minute spike tolerance) were used to define a non-wear period and 193 these periods were subtracted from daily wear time (Catellier et al., 2005). Children were included within analysis if they wore the monitor for a minimum of 9 hrs on any three days 194 195 (Mattocks et al., 2008).

PA data were classified into light (LPA), moderate (MPA), vigorous (VPA) and moderate to
vigorous PA (MVPA) intensities using three sets of intensity cutpoints: two sets of empirical
cutpoints: Evenson et al., 2008 (Ev) and Mackintosh et al., 2012 (Mack). The Mack

199 thresholds were generated from data derived from a field-based observational protocol with 200 children of the same age and from a similar geographical location as those included within 201 this study and were included to provide an additional comparison. PA was also classified using 202 individually calibrated (IC) cut points. Sedentary time was defined as ≤ 100 counts per minute 203 for all cut point sets (Fischer, Yildirim, Salmon, & Chinapaw, 2012). Individually calibrated 204 cut points were generated using the data from the VO_2 peak treadmill protocol. Froude 0.25 205 and 0.50 represent the thresholds for optimum walking speed and the transition between 206 walking and running. The average counts for the middle 90 seconds (18 epochs) of the two 207 Fr stages (Fr 0.25 and Fr 0.50) were used to represent MPA and VPA thresholds for each individual. The middle 90 seconds were selected to avoid the transitional periods between the 208 209 Fr.25 (walking) and Fr.5 (running). To examine the energy cost associated with each Fr 210 threshold metabolic equivalents were calculated for Fr.25 and Fr.5 stages using the gas analysis data (1 MET = $4.59 \text{ VO}_2 \text{ ml/kg/min}$; (Ridley & Olds, 2008) and compared to the 211 212 energy costs outlined by Harrell et al. (2005). This gas analysis data was simply used to assess 213 the MET values associated with the Fr stages for each individual.

214

215 Data analysis

216 Differences in anthropometrics, clustered risk, VO₂peak, sedentary time and PA components 217 (MPA, VPA, MVPA) were examined by sex using MANCOVA, controlling for accelerometer 218 wear time. The prevalence of those reaching \geq 60mins MVPA per day was calculated for each 219 cutpoint set. Differences in PA intensities were examined using repeated measures ANOVAs 220 by cutpoint. To investigate the association between PA components and health markers 221 (VO2peak, BMI Z-score, waist circumference, clustered cardiometabolic risk) multiple regression was employed controlling for sex, maturation, BMI and wear time. For each 222 dependent variable three multiple regression models, one for each cutpoint, were created. 223 Where BMI was used as a dependent variable it was excluded as a covariate. All analyses 224 were completed using SPSS V21.0 (SPSS Inc, IBM). Alpha was set at $P \le 0.05$. 225

226 <u>Results</u>

Unadjusted mean participant characteristics and adjusted mean anthropometric, VO₂peak,
sedentary time and PA values for boys and girls are illustrated in tables 1 and 2 respectively.
Boys were significantly less mature, had higher VO₂peak and accrued more LPA and MPA
than girls.

231

232 TABLE 1 ABOUT HERE

- 233 TABLE 2 ABOUT HERE
- 234

The IC cut points ranged from 1234-4476 counts per minute for MPA and 3192-9357 counts 235 236 per minute for VPA The mean oxygen consumption (VO₂ ml·kg⁻¹·min⁻¹) and MET values (1 MET = $4.59 \text{ VO}_2 \text{ ml} \cdot \text{kg}^{-1} \cdot \text{min}^{-1}$) achieved during the treadmill stages Fr0.25 and Fr.5 were 237 238 20.1ml·kg⁻¹·min⁻¹ (SD = 4.2 ml·kg⁻¹·min⁻¹), 4.4 METs and 31.2 ml·kg⁻¹·min⁻¹ (SD = 7.4 ml·kg⁻¹·min⁻¹) ¹·min⁻¹) 6.8 METs respectively. These values are proximal to those commonly used to 239 240 represent MPA (\geq 4 METS+) and VPA (\geq 6 METS) in the PA literature. Data from Harrell et 241 al (2005) calculated for children aged 8-12 years confirm that participants were working at an 242 intensity approximately equivalent to moderate intensity during Fr 0.25 (Harrell et al. (2005) values: $VO_2 = 18.3 \text{ ml}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$) and approaching high intensity activity during Fr 0.50 243 244 (Harrell et al. (2005) values: $VO_2 = 38.5 \text{ ml} \cdot \text{kg}^{-1} \cdot \text{min}^{-1}$).

245

The proportion of children meeting ≥60mins/day MVPA varied depending on the cutpoints
used. According to the Mack cutpoints 56% met ≥60mins/day MVPA, whereas 49% and 37%
achieved 60mins according to Ev and IC cutpoints respectively.

The results of the repeated measures ANOVAs between cutpoint sets can be viewed in Table 3. Significantly higher MPA was reported using the Mack (MPA = 51.6 mins/day) cutpoints in comparison to the Ev (MPA = 38.4 mins/day) and IC cutpoints (MPA = 44.3 mins/day). Ev cutpoints (VPA = 25.9 mins/day) recorded higher VPA than Mack (17.8 mins/day) and IC (13.0 mins/day). Significantly less LPA was observed using the Mack (195 mins/day)
cutpoints in comparison to IC (209 mins/day) and Ev (200.3 mins/day), the difference between
IC and Ev LPA was also statistically significant.

256

257 TABLE 3 ABOUT HERE

258

Results of multiple regression found that IC LPA and MPA were significant predictors of 259 VO_2 peak (R² for the model = 0.55, LPA: standardised beta = 0.32, t= 3.24, p = 0.002, MPA: 260 standardized beta = 0.27, t = 2.57, p = 0.013) IC MPA was also a significant predictor for BMI 261 Z-score (R^2 for the model = 0.31, standardised beta =- 0.35, t = -2.96, p = 0.004). Ev VPA was 262 a significant predictor for BMI Z-score (R^2 for the model = 0.32, standardized beta = -0.33, t 263 264 = -2.59, p = 0.012), Mack data were not significant predictors for any health variables, however Mack VPA approached statistical significance as a predictor for BMI Z-score (R² for 265 266 the model = 0.29, VPA standardized beta = -0.23, t = -1.98, p = 0.052). No significant PA-267 clustered risk score associations were observed irrespective of cut point set used.

268

269 Discussion

270 The aims of this study were to compare children's physical activity (PA) levels, the prevalence 271 of children meeting current guidelines of 60 minutes of daily moderate to vigorous PA 272 (MVPA), and PA-health associations using individually calibrated Fr (IC) and empirical 273 accelerometer cutpoints. In this study the proportion of the cohort meeting current guidelines 274 for daily MVPA ranged from 37% - 56% depending upon the cut point used. A number of 275 studies have shown differences in PA prevalence depending on the choice of cut points used 276 to analyse data (Hislop et al., 2012; Reilly et al., 2008). For example, a review by Ekelund et 277 al. (2011) highlighted that the reported prevalence of children and young people meeting current PA guidelines ranged across six studies from 1% and 100%, with authors suggesting 278 that the variability could be largely attributed to the different intensity cutpoints used between 279 280 studies. At a 4 MET intensity (approximate to MPA) recommended counts per minute have

ranged widely from 1400 to 3600 (Cain et al., 2013). In this study, IC cut points derived from
the treadmill-based protocol ranged from 1234-4476 for MPA, with a mean MET value of 4.4
(range 2.49-7.04 METs) and 3192-9357 for VPA (mean MET value of 6.8, range 3.83-12.33
METs). This demonstrates the substantial variation that exists in the biomechanical efficiency
of movement (e.g. stride length, stride pattern) between children of a similar age completing
the same activity, and provides support for the use of IC cut points that take account of
individual differences.

288

289 This study also revealed significant differences in the classification of LPA, MPA and VPA depending upon the cutpoint used. The discrepancies in the classification of PA intensities 290 291 observed may be in part due to the differing methods used to define each of the cutpoint 292 thresholds. For example, the IC cut points used in this study were derived from the application 293 of biomechanical theory, which is in contrast to the empirical cut points derived from 294 laboratory based (Ev) or field-based (Mack) energy expenditure. Although it is well 295 documented that the application of different cut points results in differences in estimates of 296 activity intensity (Trost et al., 2011), to date, none of the published papers have compared the 297 classification of activity intensity between IC and empirical cut points. Our findings suggest 298 that researchers should be cautious about the universal application of cut points which fail to 299 account for individual differences between participants, particularly with evidence suggesting 300 wide variations in step counts between children when engaging in PA at equivalent intensities 301 (Rowlands, 2007). The application of more specific cutpoints may provide an opportunity to 302 reduce sample size requirements within studies due to better estimates of primary outcome 303 measures.

304

305 Despite calls for raw data processing techniques to remove the reliance on proprietary counts 306 based data, this approach still requires the use of cutpoints or acceleration thresholds to 307 classify raw acceleration signals, therefore the findings of this study apply in the raw data 308 analysis context. To remove the requirement of cutpoints, pattern recognition or machine

309 learning approaches to classify accelerometer data have been proposed Despite the potential 310 utility of this approach the majority of PA research conducted to date using accelerometers 311 still utilises proprietary counts data and apply group level thresholds to the data, therefore the 312 IC approach proposed within this paper is recommended. The range in prevalence and 313 classification of PA resulting from the application of different cut points underscores the need 314 for a consensus on accelerometer thresholds to quantify PA intensity. The array of thresholds 315 used by researchers makes comparison between studies problematic, leading to conflicting 316 conclusions (Hislop et al., 2012). The inconsistent use of these thresholds is also a major issue 317 when attempting to quantify the prevalence of inactivity (Ekelund et al., 2011), has impacted upon PA policy making for children (Bailey et al., 2013) and the relationships between PA 318 319 and health outcomes (Bailey et al., 2013). For example, in their comparison of three published 320 thresholds (Chu, McManus, & Yu, 2007; Rowlands, Thomas, Eston, & Topping, 2004; Vanhelst et al., 2010) Bailey et al. (2013) reported a range of different associations between 321 322 PA and health outcomes such as blood pressure, waist circumference, cardiorespiratory fitness 323 and metabolic markers such as glucose and triglycerides. In our study, although the estimates 324 of the intensity of PA differed according to thresholds used, relationships detected with the 325 clustered cardiometabolic risk score were consistent. However, the IC cut points had the 326 strongest associations with VO₂peak, an important independent predictor of cardiometabolic 327 risk (Andersen et al., 2011). This may be due to the methods utilised to create the IC cutpoints 328 and the treadmill-based VO₂peak protocol. Whether the approach of individually calibrating 329 PA thresholds according to limb length is as effective at predicting energy expenditure across 330 a range of different activities warrants further investigation. Furthermore, the empirical 331 cutpoints used in this study were created using field-based protocols that included a range of typical daily activities. The differences described between these methods and the IC approach 332 333 may relate to the protocols used to generate the cutpoints, rather than the accuracy of the 334 cutpoints per se.

336 When compared to the use of a portable metabolic unit (Ev) or PA observation (Mack), it appears that the treadmill-based calibration protocol used in this study was more effective in 337 338 accounting for individual differences in biomechanical efficiency of movement (e.g limb 339 length, stride length/frequency) by matching the accelerometer counts to changes in speed and 340 resulting PA intensity. Whereas in previous research, the observed relationships between CRF and PA in children have been weaker than expected, for example weak-moderate standardized 341 regression coefficients (0.14-0.33) between aerobic fitness and PA have been 342 343 reported(Kristensen et al., 2010), the present findings suggest that IC cut points highlight stronger associations between children's PA and CRF than are often reported. Moreover, there 344 is a growing body of evidence that links CRF to cardiometabolic disease risk in children 345 (Anderssen et al., 2007) (Boddy et al., 2014). Therefore, our findings have important 346 347 implications for researchers investigating the associations between activity status, CRF and 348 health, and practitioners referring inactive individuals for lifestyle intervention.

349

350 Strengths and Limitations: This is the first study to examine differences in reported PA, and 351 PA-health relationships between Fr IC and empirically derived cut points. The generic 352 cutpoint method is less time consuming than completing laboratory calibration studies 353 involving multiple activities and portable calorimetry or observation. Despite this, the 354 individual calibration approach used within this study does not take into account movement 355 patterns other than walking and running, and although the majority of children's activity is 356 ambulatory the method may not accurately classify other types of movements completed by 357 children. The method also did not merge the VO₂ data from the fitness assessment that would have provided energy expenditure data. This was purposeful to allow the examination of the 358 thresholds based on the Fr number alone, rather than a more complex hybrid threshold 359 approach. An evolution of this method could be proposed that utilises VO₂ data to examine 360 whether the precision of the thresholds is improved, however this was beyond the scope and 361 362 aims of the current study. In addition, the Fr number could result in non-ecological walking 363 patterns which are not representative of 'usual' walking speeds. It is important to note that 364 maturational factors may influence metabolic efficiency and therefore energy expenditure 365 within this population. The influence of maturation on energy expenditure was not explored 366 within this study, mainly because of the repeated measures nature of the analysis when 367 comparing thresholds, however warrants consideration when working with populations within 368 this age range. This study used a range of established and emerging risk factors to provide a 369 robust estimate of cardiometabolic risk. However, the participants involved in this study were 370 healthy children, which may account for the lack of associations observed between the PA 371 data and cardiometabolic risk scores. Stronger PA-health associations may be apparent in a 372 population exhibiting greater cardiometabolic risk. The treadmill measure of VO₂peak is 373 considered the reference standard, though standardised protocols were used, data were taken 374 using different gas analysis systems in Liverpool and Ulster (Oxycon Pro and COSMED) which may influence comparability between the VO₂peak estimates. 375

376

377 Conclusion

378 This study has demonstrated that the application of different intensity thresholds has an impact 379 when determining the proportion of children meeting current daily PA guidelines. To make accurate evidence based recommendations, a consensus on appropriate accelerometer 380 381 thresholds for quantifying PA intensity is needed. IC cut points provide evidence of a stronger 382 association between children's PA and CRF than is often reported. This finding has important 383 implications for researchers and practitioners investigating the associations between activity 384 status, CRF and health and referring inactive individuals for lifestyle intervention. Additional 385 research is needed with larger cohorts to fully examine the potential of using IC cut points to 386 classify children's PA.

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Variable	Boys (n = 32)	Girls $(n = 43)$
Age (years)	10.47 [0.57]	10.5 [0.75]
Maturation offset (years)	-2.64 [0.71]	-0.95 [0.68]
BMI Z-score	0.65 [0.86]	0.53 [1.24]
Waist circumference (cm)	64.4 [6.4]	65.7 [9.57]
VO ₂ peak (ml/kg/min)	45.55 [9.71]	40.81 [8.7]
Diastolic BP (mmHg)	63 [6.2]	62.1 [6.7]
Systolic BP (mmHg)	103.6 [11.9]	102.3 [12.1]
FMD %	8.39 [3.24]	8.54 [4.26]
C-Reactive Protein (mg/L)	0.38 [0.29]	0.94 [1.32]
Triglycerides (mmol/L)	0.64 [0.2]	0.78 [0.28]
Cholesterol (mmol/L)	4.17 [0.67]	4.21 [0.54]
HDL-C (mmol/L)	1.59 [0.31]	1.49 [0.38]
Glucose (mmol/L)	4.71 [0.34]	4.63 [0.3]
Adiponectin (µg/mL)	10.58 [5.4]	11.14 [6.78]
Clustered CM risk	0.18 [4.01]	-0.38 [3.71]
Sedentary Time (mins/day)	440.4 [41]	458.8 [41.2]

494 <u>Table 1. Raw mean [SE] participant characteristics by sex</u>

IC MPA (mins/day) IC VPA (mins/day)	52.4 [36.3] 14.2 [9.7]	38.3 [24.7] 12.2 [17.7]
IC VPA (mins/day)	14.2 [9.7]	12.2 [17.7]
C MVPA (mins/day)	66.6 [37.5]	50.4 [31.9]
Accelerometer wear time	716.8 [49.2]	716.6 [116.9]

Variable	Boys	Girls	P value
Age (years)	10.45 [.13]	10.49 [.12]	.833
Maturation offset (years)	-2.63 [.14]	-0.89 [.13]	< 0.001
BMI Z-score	0.62 [.21]	0.70 [.18]	.776
Waist circumference (cm)	64.0 [1.6]	67.0 [1.5]	.173
VO ₂ peak (ml/kg/min)	46.39 [1.61]	39.28 [1.44]	.002
Clustered CM risk	-0.047 [.75]	-0.282 [.67]	.816
Sedentary Time (mins/day)	438.6 [7.7]	457.8 [6.9]	.069
IC LPA (mins/day)	221.5 [7.4]	196.3 [6.6]	.014
IC MPA (mins/day)	53.5 [5.9]	37.4 [5.3]	.047
IC VPA (mins/day)	14.0 [3.1]	12.6 [2.8]	.741
IC MVPA (mins/day)	67.5 [7.0]	50.1 [6.2]	.066

496 <u>Table 2. Mean values [SE] for boys and girls adjusted for wear time (MANCOVA</u>

Activity	Individual	ly Calibrated	Mackinto	sh et al. 2012	Evenson	et al., 2008
Component	(IC) Minu	tes/day	(Mack) M	linutes/day	(Ev) Mir	utes/day
	Mean	SE	Mean	SE	Mean	SE
VPA	13.0*	1.7	17.8†	1.7	25.9	2.0
MPA	44.3^^	3.6	51.6^	2.4	38.4	1.8
LPA	209.0*	4.1	195.0 [†]	3.7	200.3	3.8
501						

Table 3. Adjusted mean [SE] physical activity across the three cutpoint sets

502	[*] Ev > IC (p < 0.001), ["] Mack > IC (p < 0.001), [†] Ev > Mack (p < 0.001), [^] Mack > IC
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(p = 0.005) ^Mack > Ev (p < 0.001), [‡]IC > Mack (p < 0.01) and IC > Ev (p = 0.006).