

**Title**

Comparability of published cut-points for the assessment of physical activity: Implications for data harmonization

**Running head:**

Effect of cut-points on physical activity

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## **Abstract**

This study aimed to compare estimations of sedentary time (SED) and time spent in physical activity (PA) intensities in children with overweight/obesity across different age-appropriate cut-points based on different body-worn attachment sites and acceleration metrics. A total of 104 overweight/obese children (10.1±1.1 years old, 43 girls) concurrently wore ActiGraph GT3X+ accelerometers on their right hip and non-dominant wrist for 7 days (24 hours). Euclidean Norm Minus One *g* (ENMO) and activity counts from both vertical axis (VACounts) and vector magnitude (VMCounts) were derived. We calculated estimates of SED and light, moderate, vigorous, and moderate-to-vigorous (MVPA) intensity PA using different published cut-points for children. The prevalence of children meeting the recommended 60 min/day of MVPA was calculated. The time spent in SED and the different PA intensities largely differed across cut-points based on different attachment sites and acceleration metrics (i.e., SED = 11-252 min/day; light PA = 10-217 min/day; moderate PA = 1-48 min/day; vigorous PA = 1-35 min/day; MVPA = 4-66 min/day). Consequently, the prevalence of children meeting the recommended 60 min/day of MVPA varied from 8% to 96% of the study sample. The present study provides a comprehensive comparison between available cut-points for different attachment and acceleration metrics in children. Furthermore, our data clearly show that it is not possible (and probably will never be) to know the prevalence of meeting the PA guidelines based on accelerometer data since apparent differences range from almost zero to nearly everyone meeting the guidelines.

**Keywords (3-8):** Activity monitor; exercise; sedentary lifestyle; lifestyle behaviors; adolescent; youth.

## 2 **1. Introduction**

3 Accurate and objective estimations of daily sedentary time (SED) and physical activity (PA) are important  
4 to estimate the prevalence of populations meeting the current PA guidelines, to assess the success of  
5 interventions aiming to increase PA in specific populations, to explore population activity trends, and to  
6 quantify the dose-response impact of SED and PA on health <sup>1</sup>. Accelerometers are feasible tools to  
7 objectively assess SED and PA in large-scale studies, but their utilization requires standardized data  
8 collection (e.g., attachment site) and processing criteria (e.g., how to filter the raw accelerations), both  
9 demonstrating a high potential to affect the estimation of PA <sup>2</sup>. Additionally, protocols and methods vary  
10 largely across studies which aims to develop cut-points (e.g., differences in the exercise protocols or the  
11 measurement of energy expenditure), resulting in differences in the identification and application of cut-  
12 points, i.e., intensity thresholds for SED and PA intensity classification. Since SED refers to any waking  
13 behavior in a reclining posture with requires low related energy expenditure <sup>3</sup>, it is important to note that  
14 SED estimations based on cut-points are limited because they are not able to detect changes in posture.  
15 Many authors have called for a harmonization of data collection, processing criteria, and selection of cut-  
16 points to assess SED and PA in order to gain comparability between studies <sup>2,4,5</sup>. This harmonization would  
17 be of special interest to compare data across studies, especially when the populations assessed are similar.  
18 To date, such harmonization and consensus is not available.

19 Data collection decisions include selecting a device, the body attachment site (i.e., hip or wrist in the  
20 majority of studies) and the sampling frequency for the recording (usually between 30-100 Hz) <sup>2</sup>. The  
21 traditional hip attachment site is being replaced with a wrist location by some consumer-grade  
22 manufacturers (e.g., FitBit, Polar, Garmin, or Up) and by large-scale studies, such as the US National  
23 Health and Nutrition Examination Survey (NHANES) and the UK Biobank. This strategy was undertaken  
24 as an effort to obtain a higher wear compliance <sup>2,6,7</sup>. Both hip and wrist attachment sites have been validated  
25 for classifying PA intensities <sup>2,8-10</sup>, and are potentially able to assess energy expenditure during free-living  
26 conditions in different populations <sup>11,12</sup>, yet due to differences in the protocols used in cut-point validation  
27 studies it is unknown how well measures from the hip and wrist compare to each other.

28 The main purpose of processing criteria is to get a clean estimate of body accelerations by removing gravity  
29 acceleration and noise from the acceleration signal. The first commercially available accelerometers  
30 coerced researchers into using the manufacturer's activity counts (i.e., accelerations due to body  
31 movement) from the vertical axis (VACounts) or vector magnitude (VMCounts) derived from proprietary  
32 algorithms. These activity counts were hardly comparable between devices, or even between different  
33 models from the same manufacturer <sup>13,14</sup>. However, contemporary accelerometers are capable of storing  
34 high-frequency raw accelerations, which are highly comparable between frequently used research-grade  
35 devices (i.e., ActiGraph, GENEActiv, and Axivity) <sup>15</sup>. In the last five years, researchers have published  
36 open source methods to process raw accelerations in order to obtain alternative acceleration metrics to  
37 activity counts <sup>16,17</sup>. Euclidean Norm of raw accelerations Minus One *g* (ENMO) is now widely used and  
38 has shown a high agreement between brands <sup>15,18</sup>, facilitating data harmonization across studies.

39 As the process of harmonizing data collection and processing criteria proceeds, it is important to study  
40 how different body attachment sites, acceleration metrics, and cut-points affect the final estimations of  
41 SED and PA intensities. Rowlands et al. reported a moderate agreement between moderate-to-vigorous  
42 PA (MVPA) estimates derived using different cut-points based on ENMO from wrist accelerations and  
43 classical activity counts thresholds based on hip-worn devices <sup>19</sup>. In contrast, other studies comparing cut-  
44 points developed independently for different attachment sites and acceleration metrics have reported large  
45 differences across MVPA estimates in adolescents <sup>4</sup> and adults <sup>5</sup>. Although there is an increasing interest  
46 in the study of SED and light intensity PA <sup>20</sup>, previous studies have only focused on MVPA.

47 Therefore, there is a need to better understand how data collection, processing criteria, and cut-points  
48 influence estimations of SED and PA in different populations, including children and those classified as  
49 overweight/obese. Thus, this study aimed to examine how cut-points relative to different attachment sites  
50 and acceleration metrics affect the final estimations of SED and PA in children with overweight/obesity.

## 51 **2. Methods**

52 The present cross-sectional study analyzed data from the baseline assessment of the ActiveBrains Project  
53 (<http://profith.ugr.es/activebrains>). A detailed description of the study design and methods has been

54 published elsewhere <sup>21</sup>. Briefly, ActiveBrains is a randomized controlled trial intended to examine the  
55 effect of a 20-week PA intervention on brain structure, function, cognitive performance, academic  
56 achievement, and physical and mental health outcomes in overweight/obese children <sup>21</sup>. A total of 110  
57 overweight/obese children (classified based on the World Obesity Federation cut-points <sup>22,23</sup>) were  
58 recruited from Granada (Spain). A final sample of 104 children ( $10.1 \pm 1.1$  years of age, 41% girls) met  
59 the accelerometry inclusion criteria (more details below). The data were collected between November 2014  
60 and February 2016. We informed the parents or legal guardians about the purpose of the study, and we  
61 obtained written informed parental consent. The ActiveBrains project was approved by the Human  
62 Research Ethics Committee of the University of Granada, and was registered as a clinical trial  
63 (NCT02295072, <http://clinicaltrials.gov>).

64 The participants' anthropometry, SED, and PA were assessed as part of the protocol of the ActiveBrains  
65 project <sup>21</sup>. Briefly, we measured the body weight and height to the nearest 0.1 kg and 0.1 cm using an  
66 electronic scale (SECA 861, Hamburg, Germany) and a precision stadiometer (SECA 225, Hamburg,  
67 Germany), respectively. Body mass index (BMI) was calculated as  $\text{kg/m}^2$ . The participants were also  
68 required to concurrently wear two accelerometers (ActiGraph GT3X+, Pensacola, FL, USA) for 7  
69 complete days (24 hours): one on the right hip and the other on the non-dominant wrist. The participants  
70 were instructed to wear the accelerometers as many hours as possible and to remove them only for water  
71 activities (i.e., shower or swimming), and both at the same time. Concomitantly, the participants reported  
72 the time they went to bed and woke-up in a diary log throughout the study.

73 ActiGraph GT3X+ is a triaxial accelerometer with a dynamic range of  $\pm 6$  G. Both hip- and wrist-worn  
74 accelerometers were initialized to capture and store accelerations at 100 Hz. The raw accelerations were  
75 then downloaded and converted to “.csv” format using ActiLife v.6.13.3 (ActiGraph, Pensacola, FL, USA).  
76 Raw “.csv” files were imported to R software (v. 3.1.2, <https://www.cran.r-project.org/>) and processed  
77 using the GGIR package (v. 1.5-12, <https://cran.r-project.org/web/packages/GGIR/>). They were also  
78 imported and processed in the ActiLife software (ActiGraph, Pensacola, FL, USA) to obtain VMCounts  
79 and VACounts using the normal filter developed by ActiGraph. The processing methods involved: 1)  
80 Auto-calibration of the data according to the local gravity <sup>24</sup>. 2) Detection of the non-wear time based on

81 the raw acceleration of the three axes <sup>16</sup>. Briefly, each 15-min block was classified as non-wear time if the  
82 standard deviation of 2 out of the 3 axes was lower than 13 mg during the surrounding 60-min moving  
83 window, or if the value range for 2 out of the 3 axes was lower than 50 mg. 3) Detection of sustained  
84 abnormal high accelerations, i.e., higher than 5.5 g. 4) Calculation of the Euclidean Norm Minus One  
85 (ENMO) as ( $\sim 9.8 \text{ m/s}^2$ ) with negative values rounded to zero. 5) Importation of the VMCounts and  
86 VACounts “.csv” files to R to follow the same processing criteria than ENMO. 6) Imputation of detected  
87 non-wear time and abnormal high accelerations by means of the acceleration for the rest of the recording  
88 period during the same time interval than the affected periods. 7) Identification of waking and sleeping  
89 hours using an automatized algorithm guided by the times reported by the participants <sup>25</sup>. Waking and  
90 sleeping hours were detected using data from the non-dominant wrist and detected times were then  
91 matched to the right hip data for each participant. And, 8) Estimation of SED and PA intensities using  
92 different age-appropriate cut-points for ENMO, VMCounts, and VACounts as detailed in **Table 1**.

93 Mean daily SED and PA intensity levels were then calculated as: (mean of available weekdays\*5 + mean  
94 of available weekend days\*2) / 7. The participants were excluded from the analyses if they recorded less  
95 than 4 valid days (i.e.,  $\geq 16$  hours/day), including at least 1 weekend day. Out of the 110 participants, 4  
96 children recorded less than 4 days of valid wearing time, 1 accelerometer attached to the non-dominant  
97 wrist malfunctioned, and 1 participant was excluded for having mean acceleration values during nights  
98 between 6-9 standard deviations above the group mean. Thus, a final sample of 104 participants was  
99 included in the present study.

100 Descriptive statistics were calculated as means and standard deviations. The time estimates of SED, light,  
101 moderate, vigorous intensity PA, and MVPA were compared between each pair of estimations (i.e.,  
102 estimations from each pair of cut-points) using repeated measures analysis of variance (ANOVA).  
103 Additionally, we inspected the distributions of the time spent in MVPA and the prevalence of the study  
104 sample meeting the PA guidelines (i.e., at least 60 min/day of MVPA) <sup>26</sup> using different cut-points. All  
105 analyses were performed in R. Overall, the significance level was set at  $p < 0.05$  for all the analyses;  
106 however, in order to account for multiple comparisons, significant differences at  $p < 0.01$  were interpreted  
107 as statistically meaningful.

### 108 3. Results

109 The anthropometric characteristics, the time spent in SED, and the various PA intensities (calculated using  
110 the different cut-points) are reported in **Table 2**.

111 The comparisons between SED and PA intensities estimated from the different cut-points are graphically  
112 presented in **Figure 1**. The differences expressed in min/day between different cut-point estimates are  
113 shown in **Table 3**. Nearly every pairwise comparison was significantly different (all  $p < 0.05$ ) (exceptions  
114 are shown in Table 3). Overall, the various mean daily estimations differed between 11-252 min/day for  
115 SED, 10-217 min/day for light intensity PA, 1-48 min/day for moderate intensity PA, 1-35 min/day for  
116 vigorous intensity PA, and 4-66 min/day for MVPA.

117 **Figure 2** presents the time distributions spent in MVPA for the different cut-points examined. Overall,  
118 this figure shows that cut-points based on VMCounts produced higher MVPA time compared to those  
119 estimations based on ENMO or VACounts, independently of the attachment site (as reported in Table 3).

120 **Figure 3** shows that the sample prevalence meeting the recommended 60 min/day of MVPA per day  
121 ranged from 8% to 96% depending on the cut-points applied to the data. Overall, the prevalence of meeting  
122 the PA guidelines was higher for boys than for girls using all cut-points except for the Chandler et al. <sup>9</sup>  
123 cut-points (i.e., 90% of the boys versus 95% of the girls met the PA guidelines, accordingly).

### 124 4. Discussion

125 The primary purpose of this study was to provide a clear picture of which cut-points are more and less  
126 comparable in free-living conditions in children with overweight/obesity, including traditional (e.g.,  
127 Evenson cut-points based on VACounts <sup>27</sup>) and recently developed (e.g., Hildebrand cut-points based on  
128 ENMO <sup>8,28</sup>, Romanzini <sup>10</sup> and Chandler <sup>9</sup> cut-points based on VMCounts) cut-points, and when the  
129 accelerometer was attached to the hip and wrist. Contrary to what could have been expected, all cut-points  
130 based on VMCounts produced significantly higher estimations of time spent in MVPA than ENMO and  
131 VACounts cut-points, regardless of the attachment site. To our knowledge, this is the first study  
132 investigating differences across accelerometer-based estimations of SED and PA intensities using a

133 complete set of available cut-points, running from the most traditionally used cut-points for VACounts  
134 detected from a hip attachment, i.e., the Evenson et al.<sup>27</sup> cut-points, to the newly developed cut-points for  
135 ENMO<sup>8,28</sup> and VMCOUNTS<sup>9,10,29</sup> from both hip and non-dominant wrist attachments.

136 Since the selection of the different data collection and processing criteria are known to affect SED and PA  
137 intensity estimations<sup>2</sup>, we applied cut-points specifically developed for the two different attachment sites  
138 for use in children. We also followed the same processing criteria (i.e., same acceleration metric and epoch  
139 length) as originally used in validation studies. In agreement with recent studies<sup>5,30</sup>, our results confirm  
140 non-comparable estimates of the time spent in MVPA when using different data collection and processing  
141 criteria. However, the present study expands upon this knowledge by additionally comparing estimates of  
142 SED and a complete range of PA intensities in a sample of overweight/obese children. Each of these  
143 metrics also displayed non-comparable estimates with large differences between cut-points (see Table 3  
144 and Figure 1). Hildebrand et al.<sup>8,28</sup> developed two sets of cut-points in the same sample to get similar  
145 estimations of SED and PA intensities from the hip and the non-dominant wrist. In contrast, herein the  
146 estimations for SED and PA for all intensities varied greatly when using the Hildebrand et al. cut-points  
147<sup>8,28</sup> for hip and wrist. This inconsistent result agrees with the Smith et al. findings<sup>4</sup>, who reported different  
148 estimations derived from two sets of cut-points developed in the same sample and differing only in the  
149 acceleration metrics (i.e., VACounts and VMCOUNTS). Our results, together with those from Smith et al.<sup>4</sup>,  
150 confirm that cut-points from different attachment sites or different acceleration metrics that are comparable  
151 in a certain sample could largely differ in others as a result of population-specific features, which may  
152 contribute to these differences in SED and PA estimations.

153 Rowlands et al.<sup>19</sup> looked for ENMO-based cut-points from the non-dominant wrist which could replicate  
154 the traditional PA estimations from the Evenson et al.<sup>27</sup> cut-points (applied to VACounts from the hip).  
155 Specifically, they reported moderate agreement (intraclass correlation coefficient -ICC- of 0.76) and 2  
156 min/day more of MVPA when applying a cut-point of 250 mg for ENMO from wrist compared to the  
157 Evenson et al.<sup>27</sup> cut-point. Accordingly, we used a lower cut-point for MVPA for ENMO wrist (i.e., 200  
158 mg – validated by Hildebrand et al.<sup>8</sup>) and detected 15 min/day more of MVPA from ENMO wrist  
159 compared with the Evenson et al.<sup>27</sup> cut-point on hip. Thus, higher values of MVPA can be expected when



160 using the cut-point by Hildebrand et al.<sup>8</sup> for ENMO wrist compared to the MVPA threshold by Evenson  
161 et al.<sup>27</sup> for VACounts hip. A more comparable threshold to identify MVPA from ENMO wrist could be  
162 250 mg<sup>19</sup>.

163 Taking these findings into consideration, the selection of cut-points to estimate PA intensities with  
164 accelerometers is a major obstacle to overcome in objective monitoring since different cut-points could  
165 lead to wildly discrepant conclusions. For example, in our sample, the prevalence of boys meeting the 60  
166 min/day of MVPA was higher than that for girls for all the cut-points except for the Chandler et al.<sup>9</sup> cut-  
167 points, for which the prevalence was higher in girls than in boys, i.e. 95% vs. 90%. Likewise, Figure 3  
168 shows large differences in the prevalence of our sample meeting the PA guidelines (i.e., from 8% to 96%),  
169 so the fundamental query regarding the prevalence of the population achieving healthful levels of PA is  
170 still unresolved. In this regard, Leinonen et al.<sup>31</sup> found moderate-to-high agreement between different  
171 methods to classify adults meeting the PA guidelines. It is important to consider that PA guidelines have  
172 been developed predominantly using self-reported data, thus, these estimations should be considered with  
173 caution. Several authors have proposed reporting PA using a full range of different accelerometer data  
174 collection and processing criteria until a consensus is reached<sup>4,5</sup>. However, this is not practical since  
175 reporting different and multifactor methodologies could require long explanations and high technical  
176 expertise from readers to understand these nuanced inconsistencies. Data pooling and reanalyzing raw  
177 accelerometer data may be a solution to overcome processing criteria inconsistencies and have been  
178 successfully applied (<http://www.mrc-epid.cam.ac.uk/research/studies/icad/>).

179 Although estimations of SED and PA intensities are easily understandable for the general population, we  
180 suggest that all studies using accelerometers should also report other PA indicators which are not  
181 influenced by cut-points, e.g., mean of the acceleration metric per day. As a first step to achieve this, we  
182 suggest using research-derived metrics, such as ENMO, which provides a valid estimate of free-living PA  
183 from hip and wrist attachments<sup>8,16,28</sup>. Furthermore, in contrast to traditional activity counts, such metrics  
184 enable comparability between devices<sup>15,32</sup> and they may be easier to interpret since the acceleration is  
185 expressed using a SI unit (i.e., mg). In fact, ENMO can be easily implemented in epidemiological studies  
186 using the GGIR software implemented in R (<https://cran.r-project.org/web/packages/GGIR/>). Studies

187 providing normative values for these acceleration metrics will ease the interpretation of findings in the PA  
188 measurement field. Furthermore, these normative values could help to identify acceleration values  
189 corresponding to meeting the PA guidelines, which could help to obtain a direct measure unaffected by  
190 the limitations shown by the cut-points.

191 Some limitations with this study should be acknowledged: 1) the sample analyzed herein was composed  
192 of overweight/obese children, and the results may not be generalizable to other populations; 2) the current  
193 study did not have a criterion measure for comparison that would allow us to assess the accuracy of each  
194 set of cut-points; and, 3) we used 90 accelerometers randomly placed in either hip or wrist. It could be  
195 hypothesized that the use of different accelerometer units is a source of error for the measurement.  
196 However, ActiGraph GT3X devices have shown to provide reliable estimations <sup>33</sup>, so we assume  
197 this source of error is likely to be very small in this study. Furthermore, all the estimates are  
198 derived from the same recordings, in case there is a device-related error, this error would be  
199 constant in all the estimates presented, and so, it is unlikely this will affect the findings. In contrast,  
200 this study's advantages are 1) the use of consistent data processing techniques with all the acceleration  
201 metrics (i.e., same calculation of non-wear time, waking and sleeping hours, which allow for a direct  
202 comparison between attachment sites, and acceleration metrics); and, 2) that the participants achieved high  
203 wearing time compliance, enabling the collection of a complete range of daily living accelerations.

204 In conclusion, this study shows large discrepancies in the time spent in SED and PA intensities across cut-  
205 points relative to different body attachment sites and acceleration metrics in overweight/obese children.  
206 Furthermore, we provide a comprehensive comparison between available cut-points in order to better  
207 understand which cut-points provide comparable results and which ones not. Also, our data clearly showed  
208 that it is not currently possible to know the prevalence of a population meeting the PA guidelines based on  
209 accelerometer data, with differences from nearly none to nearly everyone meeting the guidelines. Although  
210 currently elusive, data harmonization and consensus are essential to comparatively measure and  
211 communicate objectively monitored time in SED and various PA intensities across different studies.

## 212 **5. Perspectives**

213 In the present study, we provide a comprehensive overview on the comparability of available cut-points  
214 for the classification of SED, light, moderate, vigorous PA and MVPA from different accelerometer  
215 attachment sites and acceleration metrics in children. This overview allows researchers to know how  
216 comparable are their findings with other published studies, for example, it can be expected that SED  
217 derived from Hänggi et al.<sup>29</sup> and Romanzini et al.<sup>10</sup> cut-points is comparable, but large differences can also  
218 be expected for light PA classified using the same cut-points. The general belief that PA estimations from  
219 wrist-worn accelerometers provide higher values than those from hip-worn accelerometers is not supported  
220 by the current study. Other factors such as the acceleration metric used, and the cut-points themselves seem  
221 to have a higher influence in the final estimations than the accelerometer attachment site. Therefore, our  
222 results confirm previous studies and extend their findings to a different sample (overweight/obese children)  
223 and by using a complete set of published cut-points for this population. Data pooling and harmonization  
224 should be performed, as well as meta-analyses using data from cut-points validation studies to propose a  
225 consensual set of cut-points to be used in different settings/projects.

226

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**Table 1.** Children’s age-appropriate cut-points for the estimation of sedentary time (SED) and physical activity (PA) intensities.

References	Attachment	Acceleration	Epoch	SED/LPA	LPA/MPA	MPA/VPA
	site	metric	length			
Hildebrand et al. <sup>7,27</sup>	Hip	ENMO	5 sec	63 mg	143 mg	465 mg
Hildebrand et al. <sup>7,27</sup>	Wrist	ENMO	5 sec	36 mg	201 mg	707 mg
Hänggi et al. <sup>28</sup>	Hip	VMCounts	1 sec	3 c	56 c	-
Romanzini et al. <sup>9</sup>	Hip	VMCounts	15 sec	180 c	757 c	1112 c
Chandler et al. <sup>8</sup>	Wrist	VMCounts	5 sec	305 c	818 c	1969 c
Evenson et al. <sup>26</sup>	Hip	VACounts	15 sec	25 c	574 c	1003 c

ENMO: Euclidean norm minus 1 g; VMCounts: Vector magnitude counts; c: Activity counts; VACounts: Vertical axis counts; LPA: Light physical activity; MPA: moderate physical activity; VPA: Vigorous physical activity.

**Table 2.** Anthropometry, sedentary time (SED), and physical activity (PA) characteristics of participants.

	All (n=104)	Boys (n=61)	Girls (n=43)	P sex
<b>Anthropometry</b>				
Age (years)	10.1 ± 1.1	10.2 ± 1.2	9.9 ± 1.1	0.248
Weight (kg)	56.2 ± 10.8	56.8 ± 10.7	55.4 ± 11.1	0.533
Height (cm)	144.3 ± 8.3	144.9 ± 7.9	143.6 ± 8.9	0.443
BMI (kg/m <sup>2</sup> )	26.8 ± 3.5	26.9 ± 3.6	26.7 ± 3.5	0.766
<b>Wearing time during waking hours</b>				
Hip device (hours/day)	15.0 ± 0.6	15.1 ± 0.6	15.0 ± 0.6	0.569
Wrist device (hours/day)	14.8 ± 0.6	14.8 ± 0.5	14.8 ± 0.6	0.926
<b>SED (min/day)</b>				
Hip ENMO <sub>Hildebrand</sub>	817.4 ± 44.7	811.1 ± 42.9	826.3 ± 46.2	0.093
Wrist ENMO <sub>Hildebrand</sub>	565.1 ± 56.4	560.5 ± 56.3	571.6 ± 56.5	0.327
Hip VMCounts <sub>Hänggi</sub>	639.1 ± 64.8	634.4 ± 58.3	645.5 ± 73.1	0.412
Hip VMCounts <sub>Romanzini</sub>	628.3 ± 68.2	623.9 ± 65.7	634.5 ± 71.8	0.445
Wrist VMCounts <sub>Chandler</sub>	576.4 ± 53.9	577.4 ± 54.7	575.1 ± 53.3	0.828
Hip VACounts <sub>Evenson</sub>	600.6 ± 70.1	593.0 ± 69.7	611.1 ± 69.9	0.198
<b>LPA (min/day)</b>				
Hip ENMO <sub>Hildebrand</sub>	65.8 ± 15.8	68.4 ± 15.6	62.1 ± 15.5	<b>0.043</b>
Wrist ENMO <sub>Hildebrand</sub>	282.7 ± 38.5	279.3 ± 37.1	287.4 ± 40.3	0.298
Hip VMCounts <sub>Hänggi</sub>	176.9 ± 38.0	175.0 ± 33.3	179.5 ± 44.1	0.579
Hip VMCounts <sub>Romanzini</sub>	198.2 ± 41.5	193.6 ± 39.4	204.5 ± 44.0	0.197
Wrist VMCounts <sub>Chandler</sub>	239.0 ± 29.5	235.4 ± 29.2	244.0 ± 29.6	0.144
Hip VACounts <sub>Evenson</sub>	273.1 ± 52.1	276.4 ± 52.0	268.5 ± 52.5	0.452
<b>MPA (min/day)</b>				
Hip ENMO <sub>Hildebrand</sub>	32.9 ± 13.9	37.5 ± 14.7	26.5 ± 9.6	<b>&lt;0.001</b>

Wrist ENMO <small>Hildebrand</small>	47.5 ± 17.4	54.2 ± 18.4	38.1 ± 10.2	<b>&lt;0.001</b>
Hip VMCounts <small>Romanzini</small>	53.8 ± 14.4	57.9 ± 14.8	48.0 ± 11.7	<b>&lt;0.001</b>
Wrist VMCounts <small>Chandler</small>	81.2 ± 20.1	83.3 ± 22.7	78.4 ± 15.8	0.201
Hip VACounts <small>Evenson</small>	33.8 ± 11.5	37.9 ± 12.2	28.2 ± 7.4	<b>&lt;0.001</b>
<b>VPA (min/day)</b>				
Hip ENMO <small>Hildebrand</small>	3.0 ± 2.0	3.7 ± 2.1	2.1 ± 1.4	<b>&lt;0.001</b>
Wrist ENMO <small>Hildebrand</small>	7.6 ± 4.4	9.4 ± 4.5	5.0 ± 2.7	<b>&lt;0.001</b>
Hip VMCounts <small>Romanzini</small>	37.9 ± 16.1	44.2 ± 16.5	29.1 ± 10.6	<b>&lt;0.001</b>
Wrist VMCounts <small>Chandler</small>	6.2 ± 3.6	7.4 ± 3.7	4.6 ± 2.7	<b>&lt;0.001</b>
Hip VACounts <small>Evenson</small>	10.7 ± 6.7	12.4 ± 7.6	8.3 ± 4.4	<b>0.001</b>
<b>MVPA time (min/day)</b>				
Hip ENMO <small>Hildebrand</small>	36.0 ± 15.3	41.2 ± 16.1	28.6 ± 10.6	<b>&lt;0.001</b>
Wrist ENMO <small>Hildebrand</small>	55.1 ± 21.0	63.7 ± 22.0	43.1 ± 11.9	<b>&lt;0.001</b>
Hip VMCounts <small>Hänggi</small>	102.4 ± 26.8	110.6 ± 26.4	90.9 ± 23.1	<b>&lt;0.001</b>
Hip VMCounts <small>Romanzini</small>	91.7 ± 28.2	102.1 ± 28.7	77.1 ± 20.0	<b>&lt;0.001</b>
Wrist VMCounts <small>Chandler</small>	87.5 ± 22.5	90.6 ± 25.4	83.0 ± 16.9	0.071
Hip VACounts <small>Evenson</small>	44.5 ± 16.7	50.2 ± 18.1	36.6 ± 10.3	<b>&lt;0.001</b>

Data are presented as mean ± standard deviation. Statistically significant values are shown in bold.

Cut-points expressed with the body-worn attachment site, acceleration metric used and the first author of the validation study in subscripts, i.e., Hildebrand et al.<sup>7,27</sup>, Hänggi et al.<sup>28</sup>, Romanzini et al.<sup>9</sup>, Chandler et al.<sup>8</sup> and Evenson et al.<sup>26</sup>.

BMI: Body mass index; ENMO: Euclidean norm minus 1 g; VMCounts: Vector magnitude counts; VACounts: Vertical axis counts; LPA: Light physical activity; MPA: moderate physical activity; VPA: Vigorous physical activity; MVPA: Moderate-to-vigorous physical activity.

**Table 3.** T-tests for the comparison between sedentary time (SED), light, moderate, vigorous, and moderate-to-vigorous (MVPA) intensity physical activity (PA) calculated from different cut-points.

	SED (min/day)	LPA (min/day)	MPA (min/day)
	Difference (95%CI)	Difference (95%CI)	Difference (95%CI)
<b><i>Hip vs. hip</i></b>			
ENMO <sub>Hildebrand</sub> - VMCounts <sub>Hänggi</sub>	178 (163 to 194)**	-111 (-119 to -103)**	
ENMO <sub>Hildebrand</sub> - VMCounts <sub>Romanzini</sub>	189 (173 to 204)**	-132 (-141 to -124)**	-21 (-25 to -17)**
ENMO <sub>Hildebrand</sub> - VACounts <sub>Evenson</sub>	217 (201 to 233)**	-207 (-218 to -197)**	-1 (-4 to 3)
VMCounts <sub>Romanzini</sub> - VMCounts <sub>Hänggi</sub>	-11 (-29 to 8)	21 (10 to 32)**	
VMCounts <sub>Romanzini</sub> - VACounts <sub>Evenson</sub>	28 (9 to 46)*	-75 (-88 to -62)**	20 (16 to 23)**
VMCounts <sub>Hänggi</sub> - VACounts <sub>Evenson</sub>	38 (20 to 57)**	-96 (-109 to -84)**	
<b><i>Wrist vs. wrist</i></b>			
VMCounts <sub>Chandler</sub> - ENMO <sub>Hildebrand</sub>	11 (-4 to 26)	-44 (-53 to -34)**	34 (29 to 39)**
<b><i>Hip vs. wrist</i></b>			
ENMO <sub>Hildebrand</sub> - ENMO <sub>Hildebrand</sub>	252 (238 to 266)**	-217 (-225 to -209)**	-15 (-19 to -10)**
VMCounts <sub>Hänggi</sub> - VMCounts <sub>Chandler</sub>	63 (46 to 79)**	-62 (-71 to -53)**	
VMCounts <sub>Romanzini</sub> - VMCounts <sub>Chandler</sub>	52 (35 to 69)**	-41 (-51 to -31)**	-27 (-32 to -23)**
ENMO <sub>Hildebrand</sub> - VMCounts <sub>Chandler</sub>	-241 (-255 to -227)**	-173 (-180 to -167)**	-48 (-53 to -44)**
VMCounts <sub>Hänggi</sub> - ENMO <sub>Hildebrand</sub>	74 (57 to 91)**	-106 (-116 to -95)**	
VMCounts <sub>Romanzini</sub> - ENMO <sub>Hildebrand</sub>	63 (46 to 80)**	-85 (-95 to -74)**	6 (2 to 11)*
VACounts <sub>Evenson</sub> - ENMO <sub>Hildebrand</sub>	35 (18 to 53)**	-10 (-22 to 3)	-14 (-18 to -10)**
VACounts <sub>Evenson</sub> - VMCounts <sub>Chandler</sub>	24 (7 to 41)*	34 (22 to 46)**	-47 (-52 to -43)**

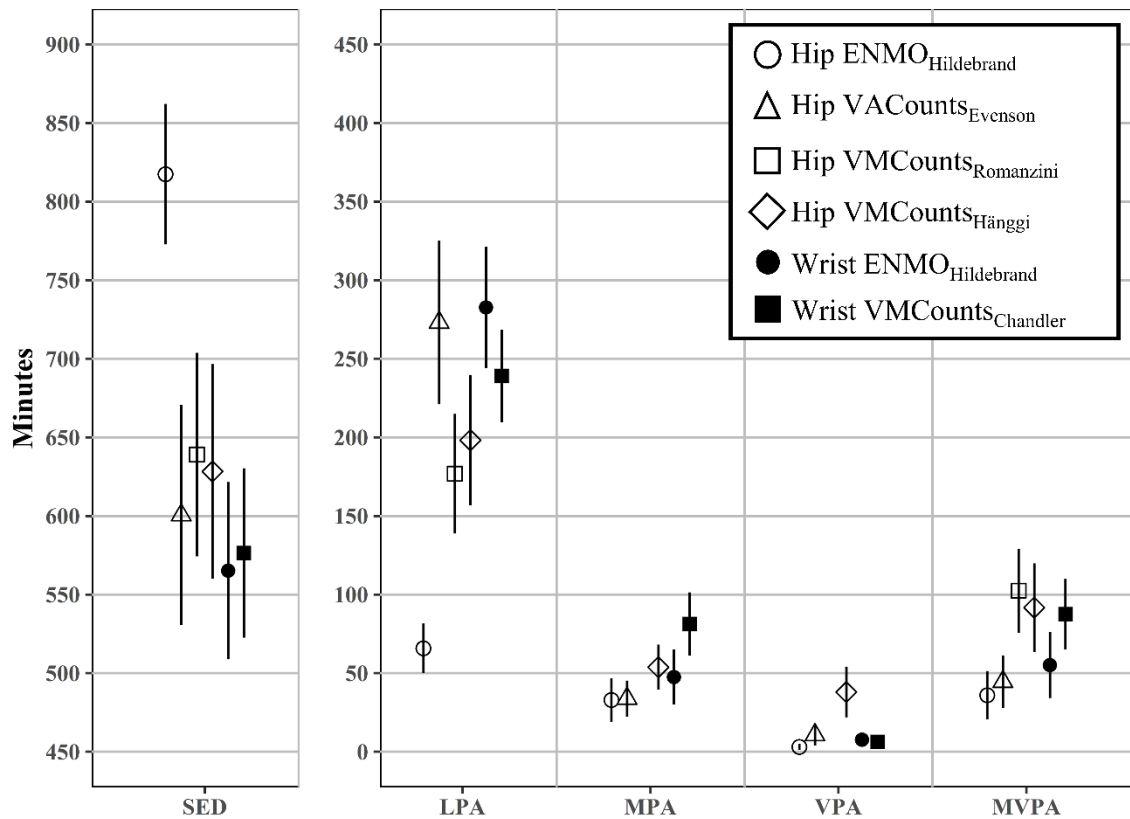
Data are presented as mean differences and 95% of confident interval.

Cut-points expressed with the body-worn attachment site, acceleration metric used and the first author of the validation study in subscripts, i.e., Hildebrand et al.<sup>7,27</sup>, Hänggi et al.<sup>28</sup>, Romanzini et al.<sup>9</sup>, Chandler et al.<sup>8</sup> and Evenson et al.<sup>26</sup>.

CI: confident interval; ENMO: Euclidean norm minus 1 g; VMCounts: Vector magnitude counts; VACounts: Vertical axis counts; LPA: Light physical activity; MPA: moderate physical activity; VPA: Vigorous physical activity; MVPA: Moderate-to-vigorous physical activity.

\*  $p < 0.05$

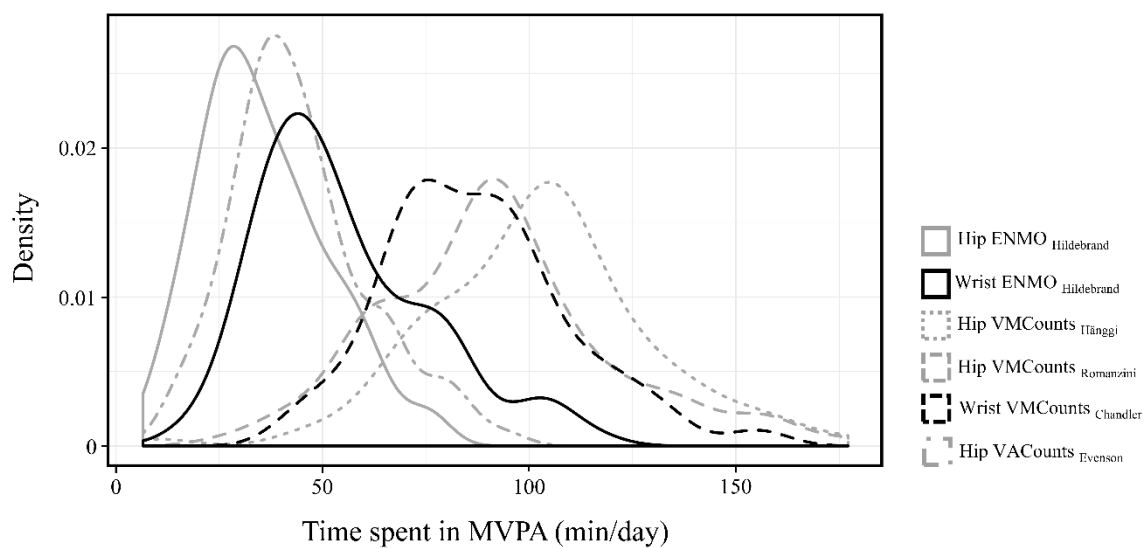
\*\*  $p < 0.01$



**Figure 1.** Mean daily time spent (min) and standard deviations (error bars) in sedentary time (SED) and physical activity (PA) considering different attachment sites and metrics.

Cut-points expressed in the legend with the body-worn attachment site, acceleration metric used and the first author of the validation study in subscripts, i.e., Hildebrand et al.<sup>7,27</sup>, Hänggi et al.<sup>28</sup>, Romanzini et al.<sup>9</sup>, Chandler et al.<sup>8</sup> and Evenson et al.<sup>26</sup>.

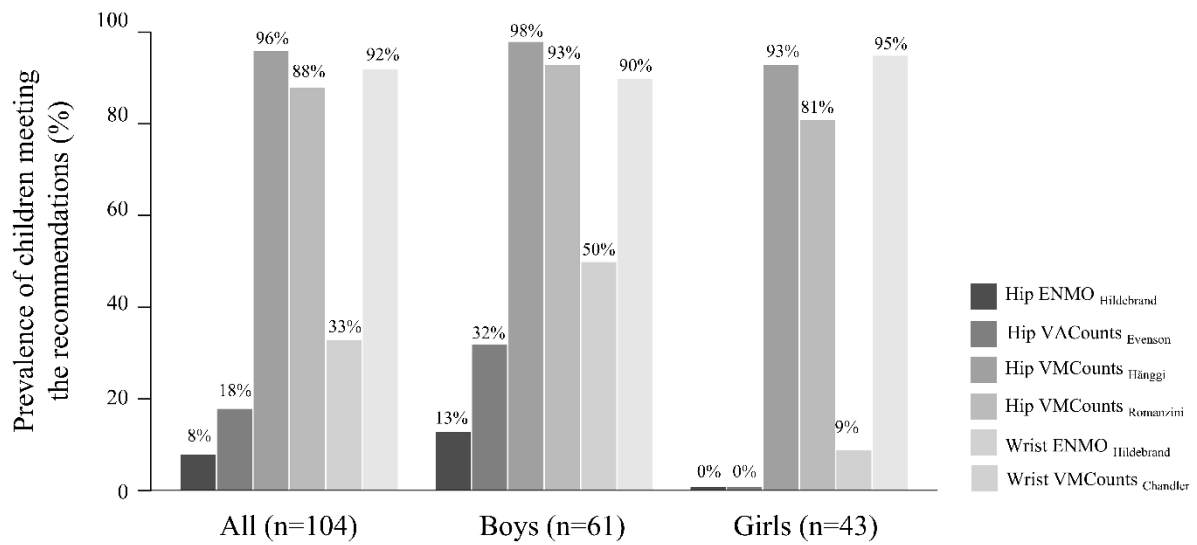
ENMO: Euclidean norm minus 1 g; VMCounts: Vector magnitude counts; VACounts: Vertical axis counts; LPA: Light physical activity; MPA: moderate physical activity; VPA: Vigorous physical activity; MVPA: Moderate-to-vigorous physical activity.



**Figure 2.** Distributions of the time spent in moderate-to-vigorous physical activity (MVPA) intensity (min/day) estimated with different cut-points.

Cut-points expressed in the legend with the body-worn attachment site, acceleration metric used and the first author of the validation study in subscripts, i.e., Hildebrand et al.<sup>7,27</sup>, Hänggi et al.<sup>28</sup>, Romanzini et al.<sup>9</sup>, Chandler et al.<sup>8</sup> and Evenson et al.<sup>26</sup>.

ENMO: Euclidean norm minus 1 g; VMCOUNTS: Vector magnitude counts; VACOUNTS: Vertical axis counts; MVPA: Moderate-to-vigorous physical activity.



**Figure 3.** Prevalence of children meeting the physical activity (PA) guidelines (i.e.,  $\geq 60$  min/day of moderate-to-vigorous physical activity -MVPA-) according to different cut-points.

Cut-points expressed in the legend with the body-worn attachment site, acceleration metric used and the first author of the validation study in subscripts, i.e., Hildebrand et al.<sup>7,27</sup>, Hänggi et al.<sup>28</sup>, Romanzini et al.<sup>9</sup>, Chandler et al.<sup>8</sup> and Evenson et al.<sup>26</sup>.

ENMO: Euclidean norm minus 1 g; VMCOUNTS: Vector magnitude counts; VACounts: Vertical axis counts.