

1 **A comparison of analytical approaches to investigate associations for accelerometry-derived**
2 **physical activity spectra with health and developmental outcomes in children**

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26 **Abstract**

27 The use of high-resolution physical activity intensity spectra obtained from accelerometry can
28 improve knowledge of associations with health and development beyond the use of traditional
29 summary measures of intensity. The aim of the present study was to compare three different
30 approaches for determining associations for spectrum descriptors of physical activity (the intensity
31 gradient, principal component analysis, and multivariate pattern analysis) with relevant outcomes in
32 children. We used two datasets including physical activity spectrum data (ActiGraph GT3X+) and 1) a
33 cardiometabolic health outcome in 841 schoolchildren and 2) a motor skill outcome in 1081
34 preschool children. We compared variance explained (R^2) and associations with the outcomes for the
35 intensity gradient (slope) across the physical activity spectra, a two-component principal component
36 model describing the physical activity variables, and multivariate pattern analysis using the intensity
37 spectra as the explanatory data matrices. Results were broadly similar for all analytical approaches.
38 Multivariate pattern analysis explained the most variance in both datasets, likely resulting from use
39 of more of the information available from the intensity spectra. Yet, volume and intensity dimensions
40 of physical activity are not easily disentangled and their relative importance may be interpreted
41 differently using different methodology.

42 **Keywords** Multivariate pattern analysis; Intensity gradient; Cardiometabolic health; Motor skills;
43 Children; Accelerometer

44

45 **Background**

46 Accelerometers capture movement across an intensity spectrum, from which summary measures of
47 time spent in different physical activity (PA) intensities, typically sedentary time (SED), light PA (LPA),
48 moderate PA (MPA), vigorous PA (VPA), and/or moderate-to-vigorous PA (MVPA), is commonly
49 derived. Although this approach is intuitively appropriate and results regarding PA levels and
50 associations with health and developmental outcomes apparently may be easily interpreted, it holds
51 important limitations. First, it requires the application of a priori defined intensity cut points, which
52 due to the lack of consistency in their application hamper comparison across studies [1]. Second,
53 limiting the description of the intensity spectrum to a few variables leads to a loss of information
54 from accelerometry [2], in particular when using linear regression analysis that cannot handle the
55 multicollinearity among the variables [3, 2].

56 Recently, two different cut point-free approaches that incorporate more detailed descriptions of the
57 PA intensity spectrum have been applied to handle these challenges in association analysis:
58 multivariate pattern analysis [4] and the intensity gradient [5]. However, the manner in which these
59 methods handle the PA intensity spectrum differs to a great extent. Aadland et al. [4] introduced
60 *multivariate pattern analysis* to analyze associations between the multicollinear explanatory PA
61 variables and cardiometabolic health in children. Multivariate pattern analysis is widely applied in
62 other fields of research with the objective of revealing patterns of important biomarkers among
63 hundreds or thousands of highly interrelated variables [6-8], and can handle completely collinear
64 explanatory variables using latent variable modelling [9, 10]. Thus, Aadland et al. [2, 4] were able to
65 determine association patterns for multiple intensity variables across the spectrum, which led to
66 improved association models compared to the use of traditional summary measures of intensity. In
67 contrast to the inclusion of multiple intensity variables in the association analysis, Rowlands et al. [5]
68 used the spectrum intensity distribution to construct the *intensity gradient*, which is a simple metric
69 that reduces an individual's intensity profile to a single variable. The intensity gradient is the slope

70 describing the curvilinear relation between time spent in lower and higher PA intensity regions (i.e.,
71 the log-log of the time-intensity curve). The intensity gradient is always negative, but is higher (i.e.,
72 the curve is flatter) the more time individuals spend in higher intensity regions [5]. The intensity
73 gradient has been shown to perform better than traditional summary measures of PA intensity (e.g.,
74 MVPA) with regard to revealing associations with health outcomes [5, 11, 12]. Thus, this approach is
75 promising given its simplicity and applicability using common statistical approaches. Description of
76 the intensity profile with a single metric also has potential for use in population comparisons and/or
77 generation of norms.

78 In addition to describing the intensity distribution in a single metric, Rowlands [5] aimed to develop
79 an intensity metric that is less dependent on the overall volume of PA. Associations between the
80 intensity gradient and overall PA level (mean acceleration) have been shown to be moderate ($r =$
81 $0.36-0.56$), which suggest the intensity gradient is more reflective of the intensity *per se* than
82 summary measures of PA intensity [5, 11, 12]. Yet, the intensity gradient and the overall PA level are
83 not independent measures of intensity and volume, respectively. Thus, research should attempt to
84 better disentangle these constructs. Principal component analysis is a well-known approach for
85 dimension-reduction of data [10], but have to the best of our knowledge not been applied to
86 describe the dimensions of intensity spectrum descriptions of PA.

87 Associations for the PA intensity spectrum with health and developmental outcomes using the
88 intensity gradient, principal component analysis, and multivariate pattern analysis have not been
89 compared. Thus, the aim of the present study was to compare associations for these three
90 approaches using two large datasets (in preschool- and schoolchildren) and two different outcomes
91 (cardiometabolic health and motor skills).

92

93 **Methods**

94 We have previously published the PA signature associated with cardiometabolic health in the *Active*
95 *Smarter Kids (ASK)* study [4, 13, 2] and the PA signature associated with motor skills in *The Sogn og*
96 *Fjordane Preschool Physical Activity Study (PRESPAS)* [14]. The aim of the present study is limited to
97 *compare* associations using multivariate pattern analysis, the intensity gradient, and principal
98 component analysis within these datasets. We refer readers to previously published descriptions of
99 sampling and children’s characteristics, study protocols, instruments, and procedures of the ASK
100 study [4, 13, 2, 15] and the PRESPAS study [16, 14] for detailed study information. Thus, we provide
101 below only a brief overview of the most relevant information to provide sufficient context to support
102 the study aim of comparing associations between these approaches.

103

104 Participants

105 The ASK study was conducted in western Norway during 2014–2015 and included 841 10-year old
106 schoolchildren providing relevant explanatory (PA) and outcome (cardiometabolic health) data [4, 13,
107 2, 15]. The PRESPAS study was conducted in western Norway during 2015–2016 and included 1081 3-
108 6-year old preschool children providing relevant explanatory (PA) and outcome (locomotor skills)
109 data [16]. Procedures and methods in both studies conform to ethical guidelines defined by the
110 World Medical Association’s Declaration of Helsinki and its subsequent revisions. The Norwegian
111 South-East Regional Committee for Medical Research Ethics and the Norwegian Centre for Research
112 Data approved the study protocols. We obtained written informed consent from each child’s parents
113 or legal guardians and from the responsible preschool and school authorities prior to all testing.

114

115 Procedures

116 *Physical activity*

117 PA was measured using the ActiGraph GT3X+ accelerometer (Pensacola, FL, USA) [17] worn at the
118 waist over seven (ASK) and 14 (PRESPAS) consecutive days, except during water activities (swimming,
119 showering) or while sleeping. Units were initialized at a sampling rate of 30 Hz and files were
120 analyzed restricted to hours 06:00 to 23:59 using 1-second epochs to capture low and high intensity
121 PA [18] using the KineSoft analytical software version 3.3.80 (KineSoft, Loughborough, UK).
122 Consecutive periods of ≥ 20 min (PRESPAS) and 60 min (ASK) of zero counts were defined as non-
123 wear time. We applied wear time requirements of ≥ 8 hours/day and ≥ 4 days/week to constitute a
124 valid measurement [19, 20].

125 We determined time (min/day) spent in PA intensities obtained from the vertical axis using
126 descriptions of 12 variables (from 0–99, 100–999, 1000–1999, ... 9000–9999, to ≥ 10000 cpm) in the
127 ASK dataset [2] and 17 variables (from 0–99, 100–999, 1000–1999, ... 14000–14999, to ≥ 15000 cpm)
128 in the PRESPAS dataset [14], to capture movement in narrow intensity intervals across the intensity
129 spectrum. These models using spectra of reduced resolutions performed similarly to previously
130 published models [2, 14] using spectra with higher resolution [21]. In the multivariate pattern
131 analysis, these spectra were included as the explanatory data matrix. We used the natural log (\ln) of
132 time to ensure comparability with the intensity gradient.

133 The concept of the intensity gradient was developed using raw acceleration data [5]. We applied the
134 theoretical premise outlined by Rowlands et al. [5] to ActiGraph count data and determined the
135 intensity gradient across the intensity spectra outlined above by calculating the slope between the \ln
136 of the intensity and \ln of the time distribution. However, while Rowlands et al. used 24-hour raw
137 acceleration data, we did not have 24-hour data and used therefore only waking time count data for
138 the analysis. Wear time was not normalized among individuals as the distribution of time (i.e., the
139 slope) is independent of the total wear time. We excluded the most extreme intensity category from
140 the calculation, since accumulated time in this larger bin caused violation of linearity of the \ln time-
141 intensity distribution. Yet, results were similar whether this bin was included or excluded. In addition

142 to the intensity gradient as a proposed measure of intensity, we included overall PA (average cpm) as
143 a measure of PA volume.

144 We included descriptive characteristics and associations with the outcomes for traditional summary
145 measures of PA intensity as supplemental material using the Evenson et al. [22, 23] intensity cut
146 points of 0–99, 100–2295, 2296–4011, and ≥ 4012 cpm to determine intensities across the spectrum
147 as SED, LPA, MPA, and VPA, respectively.

148

149 *Anthropometry*

150 In both studies, body mass was measured using an electronic scale (Seca 899, SECA GmbH, Hamburg,
151 Germany) with children wearing light clothing. Height was measured using a portable Seca 217 (SECA
152 GmbH, Hamburg, Germany). Body mass index ($\text{kg} \cdot \text{m}^{-2}$) was calculated and children were classified as
153 normal weight, overweight, or obese using the Cole et al. criteria [24].

154

155 *Metabolic health – outcome in the ASK study*

156 Aerobic fitness was measured with the Andersen intermittent running test [25]. Waist circumference
157 was measured with a Seca 201 (SECA GmbH, Hamburg, Germany) ergonomic circumference
158 measuring tape two cm over the level of the umbilicus. We calculated the waist:height ratio. Systolic
159 blood pressure were measured using the Omron HBP-1300 automated blood pressure monitor
160 (Omron Healthcare, Inc, Vernon Hills, IL, US). Serum blood samples were collected in the morning
161 after an overnight fast and analyzed for total cholesterol, triglyceride, high-density lipoprotein (HDL)
162 cholesterol, glucose, and insulin at the accredited Endocrine Laboratory of the VU Medical Center
163 (VUmc; Amsterdam, the Netherlands). We calculated the total:HDL cholesterol ratio and HOMA of
164 insulin resistance [26].

165 We calculated a composite score as the mean of six variables (systolic blood pressure, triglyceride,
166 total:HDL cholesterol ratio, HOMA of insulin resistance, waist:height ratio, and the inverse Andersen
167 test) by averaging standardized scores after adjustment for sex and age using residuals from linear
168 regression. A higher score indicates poorer cardiometabolic health. A similar approach have been
169 used previously [27].

170

171 *Motor skills – outcome in the PRESPAS study*

172 Motor skills was a sum score of three locomotor movement tasks (run, horizontal jump, hop) guided
173 by the Test of Gross Motor Development 3 test battery [28, 29]. A higher score indicates better
174 locomotor skills. Children were scored quantitatively based on whether they did or did not
175 demonstrate specific criteria for each skill based on the original scoring procedures. The criteria
176 scores were averaged for each task and the total locomotor score (minimum 0, maximum 2). The
177 score was standardized after adjustment for sex, age, body mass index, and assessor of motor skills
178 using residuals from linear regression prior to analysis.

179

180 Statistical analyses

181 *Principal component analysis.* We extracted two interpretable principal components (PCs) describing
182 the main association patterns within the explanatory data matrix including all PA variables. The first
183 component (PC 1) maximally explains the mutual variation among the variables, whereas the next
184 component (PC 2) maximally explains the most of the remaining mutual variation (etc.), with the
185 constraint that these components are mutually orthogonal (i.e., not correlated). Thus, this analysis
186 reveals the underlying association patterns of the PA variables by creating latent variables
187 maximizing explained variance among the explanatory variables. Variable loadings on each PC was
188 reported to illustrate the structure of data. On this basis, the first component was indicative of

189 volume of PA (i.e., a higher score indicates that an individual spend more time in PA and less time in
190 SED; PC_{Volume}) and the second component was indicative of intensity of PA (i.e., a higher score
191 indicates that an individual spend more time in lower intensities of PA and less time in higher
192 intensities of PA; PC_{Intensity}). Each individual's scores on these components, indicating to what degree
193 an individual scored high or low on these patterns, were used for analysis.

194 *Linear regression.* Associations between overall PA, the intensity gradient, PC_{Volume}, and PC_{Intensity}, as
195 well as associations for these explanatory variables with the outcomes (cardiometabolic health (ASK
196 dataset) and locomotor skills (PRESPAS dataset)), were determined using linear regression. For the
197 principal component analysis approach, PC_{Volume} and PC_{Intensity} were included in one joint model (since
198 variables were orthogonal). For the intensity gradient approach, overall PA and the intensity gradient
199 were analyzed using separate models due to collinearity of these variables. We determined
200 associations as standardized regression coefficients and reported the explained variance (R^2) of the
201 models for comparison of model performance.

202 *Multivariate pattern analysis.* Partial least squares (PLS) regression analysis [9] was used to
203 determine the multivariate association patterns for PA intensities (explanatory variables) with the
204 outcomes. PLS regression decomposes the explanatory variables into orthogonal linear combinations
205 (PLS components), while simultaneously maximizing the covariance with the outcome variable. Thus,
206 PLS regression is able to handle completely collinear variables through the use of latent variable
207 modelling [9]. The procedure differs from that of principal component analysis by creating
208 components that maximize the covariation with the outcome, not internally among the explanatory
209 variables. Prior to PLS regression, all variables were centered and standardized to unit variance.
210 Models were cross-validated using Monte Carlo resampling with 1000 repetitions by repeatedly and
211 randomly keeping 50% of the subjects as an external validation set when estimating the models to
212 validate the number of PLS components to be included in the model [30]. Validation is an integrated
213 part of the procedure to avoid overfitting due to inclusion of minor PLS components representing

214 noise. For each validated PLS regression model, a single predictive component was subsequently
 215 calculated by means of target projection [10, 6] to express all the predictive variance in the PA
 216 intensity spectrum related to cardiometabolic health in a single intensity vector. Selectivity ratios
 217 (SRs) with 95% CIs were obtained as the ratio of this explained predictive variance to the total
 218 variance for each PA intensity variable [31-33]. The procedure for obtaining the multivariate patterns
 219 is completely data-driven, with no assumptions on variable distributions or degree of collinearity
 220 among variables.

221 The principal component analysis and linear regression was performed using IBM SPSS v. 24 (IBM
 222 Corporation, Software Group, Somers, NY). The multivariate pattern analysis was performed using
 223 Sirius version 11.0 (Pattern Recognition Systems AS, Bergen, Norway).

224

225 **Results**

226 We included 841 schoolchildren (mean (SD) 10.2 (0.3) years old, 50% boys) and 1081 preschool
 227 children (4.7 (0.9) years old, 52% boys) who provided valid data on all relevant variables (Table 1).
 228 Children's intensity-specific PA levels are shown in Supplemental Table 1.

229 **Table 1.** Children's characteristics.

	ASK (n = 841)	PRESPAS (n = 1081)
Anthropometry		
Body mass (kg)	37.0 (8.1)	19.4 (3.3)
Height (cm)	142.9 (6.7)	109.1 (7.5)
Body mass index (kg/m ²)	18.0 (3.0)	16.2 (1.4)
Overweight and obese (%)	20.8	18.2
Waist circumference (cm)	61.9 (7.5)	-
Waist:height (ratio)	0.43 (0.05)	-
Indices of metabolic health		
Andersen test (m)	898 (103)	-
Systolic blood pressure (mmHg)	105.2 (8.4)	-
Total cholesterol (mmol/l)	4.46 (0.69)	-
HDL-cholesterol (mmol/l)	1.59 (0.35)	-
Total:HDL-cholesterol (ratio)	2.91 (0.71)	-

Triglyceride (mmol/l)	0.78 (0.38)	-
Glucose (mmol/l)	4.98 (0.32)	-
Insulin (pmol/l)	55.0 (29.8)	-
HOMA of insulin resistance (index)	1.71 (0.98)	-
Motor skills		
Locomotor skills (score)		1.3 (0.4)
Physical activity (vertical axis)		
Wear time (min/day)	795 (56)	702 (50)
Overall physical activity (cpm)	708 (272)	722 (197)
Intensity gradient		
Explained variance (%)	90 (3)	86 (3)
Constant	11.0 (0.5)	12.4 (0.7)
Slope	-1.07 (0.10)	-1.30 (0.12)

230 HDL = high-density lipoprotein; HOMA = homeostasis model assessment. All values are means (SDs) if not
231 otherwise stated.

232

233 Figure 1 shows the two extracted PCs in the two datasets. The first PCs (PC_{Volume}) in both datasets
234 explained 62.8–69.0% of the total variation among the variables and indicate that spending more
235 time in PA of any intensity is related to less time spent in SED. The second PCs ($PC_{Intensity}$) explained
236 14.4–14.8% of the remaining variation among the variables and indicate that more time spent in light
237 and moderate intensity PA is related to less time spent in vigorous PA. The total explained variances
238 of the two PCs were 77.3 and 83.8% in the ASK and PRESPAS datasets, respectively.

239 While the two PCs were orthogonal, the overall PA (cpm) and the intensity gradient were strongly
240 positively associated ($r = 0.73–0.86$) in both datasets (Table 2). Both overall PA and the intensity
241 gradient were strongly positively associated with PC_{Volume} in both datasets ($r = 0.77–0.91$), whereas
242 the intensity gradient was moderately negatively associated with $PC_{Intensity}$ ($r = -0.41–-0.40$).

243

244 **Table 2.** Bivariate correlation matrix for the explanatory variables used in the linear regression in the
245 PRESPAS dataset (upper right) and the ASK dataset (lower left and shaded).

	Overall PA	Intensity gradient	PC_{Volume}	$PC_{Intensity}$
Overall PA	-	0.86	0.86	-0.10
Intensity gradient	0.73	-	0.91	-0.40
PC_{Volume}	0.77	0.90	-	0.00

PC _{Intensity}	-0.13	-0.41	0.00	-
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246

247 Table 3 shows the associations between the PA intensity spectrum and cardiometabolic health (ASK
 248 dataset) and locomotor skills (PRESPAS dataset) using the intensity gradient and principal component
 249 analysis as determined using linear regression. Associations for traditional summary measures of PA
 250 intensity are shown in Supplemental Table 2. Due to the strong associations between overall PA and
 251 the intensity gradient, we analyzed these variables in separate models. Among all variables, the
 252 intensity gradient was the single variable that was most strongly associated with the outcomes in
 253 both datasets ($R^2 = 14.0$ and 6.1% in the ASK and PRESPAS datasets, respectively). In the ASK dataset
 254 (i.e., for cardiometabolic health), the association for the intensity gradient was considerably stronger
 255 than for overall PA, whereas the associations for these variables were rather similar in the PRESPAS
 256 dataset (i.e., for motor skills). However, in comparison with the intensity gradient, the two
 257 orthogonal PCs led to an improved model fit in both datasets ($R^2 = 17.4$ and 6.5% in the ASK and
 258 PRESPAS datasets, respectively). In the ASK dataset, both a higher volume and a higher intensity
 259 were associated with better cardiometabolic health. In contrast, only volume was significantly
 260 associated with locomotor skills in the PRESPAS dataset.

261

262 **Table 3.** Associations for the intensity gradient and principal components indicative of physical
 263 activity volume and intensity with cardiometabolic health and motor skills.

Analytic approach	Cardiometabolic health (ASK)		Motor competence (PRESPAS)	
	Coeff. (p-value)	Model R^2	Coeff. (p-value)	Model R^2
Intensity gradient				
Overall PA (cpm)	-0.18 (< .001)	3.1	0.21 (< .001)	4.4
Intensity gradient (slope)	-0.38 (< .001)	14.0	0.25 (< .001)	6.1
Principal component analysis				
PC _{Volume} (score)	-0.27 (< .001)		0.25 (< .001)	
PC _{Intensity} (score)	0.31 (< .001)	17.4	-0.05 (.083)	6.5

264

265 Figure 2 shows the multivariate association patterns between PA and cardiometabolic health (ASK
 266 dataset) and between PA and locomotor skills (PRESPAS dataset). In the ASK dataset, the strongest

267 association with cardiometabolic health was found for 7000–7999 cpm. In the PRESPAS dataset, the
268 strongest association with motor skills was found for 10000–10999 cpm. Explained variances for the
269 multivariate pattern models were 20.5% (6 PLS components) and 7.4% (2 PLS components) in the
270 ASK and PRESPAS datasets, respectively. Finally, associations for all three approaches (principal
271 component analysis, the intensity gradient, and multivariate pattern analysis) were stronger than for
272 the traditional summary measures of PA intensity, though differences were minor for motor skills.

273

274 **Discussion**

275 In the present study we used two large datasets in children to explore associations between two
276 different outcomes (cardiometabolic health and motor skills) and spectrum descriptions of PA using
277 three different approaches to handle the intensity spectrum. While the intensity gradient and
278 principal component analysis reduce the dimensions of the intensity spectrum to simpler metrics
279 prior to conducting association analysis, multivariate pattern analysis retains the full intensity
280 spectrum for analysis and interpretation. Thus, the approaches differ with regard to how much of the
281 information captured by the descriptor of the accelerometry data that is subsequently retained for
282 analysis of associations with outcomes. Consistent with these different features of the analytical
283 approaches, multivariate pattern analysis led to the best model fit, indicating that this approach
284 retains relevant information from the accelerometry data that is lost when applying the other
285 approaches. However, results were broadly consistent between all three approaches. Thus, a key
286 question, is how results from these different approaches can be interpreted in practical terms.

287 Aadland et al. have previously shown that the use of multivariate pattern analysis and the inclusion
288 of multiple variables across the intensity spectrum can increase the variance explained by PA in
289 relation to health outcomes significantly [4, 18, 13, 2]. These findings result from the high-resolution
290 descriptor capturing more of the available information from the accelerometers in combination with
291 the use of an analytical approach that allows for appropriate modelling of this information [2]. Since

292 the PA variables across the intensity spectrum are highly correlated, approaches other than multiple
293 linear regression may be needed to handle such data. However, such data have certain distributional
294 and structural features which allow for reducing the complexity of the data to simpler metrics, like
295 the intensity gradient or orthogonal PCs. If such dimension reduction methods can be demonstrated
296 to retain sufficient information in the data and provide (comparable) interpretable findings, it may
297 provide simple solutions to handle the multicollinearity of the PA intensity spectrum in association
298 analysis, which may be particularly attractive for researchers with less advanced statistical expertise.
299 Consistent with previous studies [12, 11, 5], our findings showed that the intensity gradient
300 explained more variance in outcomes compared to the traditional summary measures of PA, in
301 particular in relation to cardiometabolic outcomes. Still, association models improved further when
302 using principal component analysis, though both these approaches explained less variance than the
303 use of multivariate pattern analysis. These findings suggest dimension reduction methods to
304 construct simpler metrics of the PA intensity distribution or data structure lead to a loss of
305 information retained for association analysis compared to the use of the high-resolution intensity
306 spectrum in multivariate pattern analysis.

307 Beyond overall model performance, a crucial point that deserves attention is to which extent the
308 three models lead to similar interpretations, or whether they may lead to new knowledge of
309 associations between PA and health and developmental outcomes. Specifically, our results may
310 provide new perspectives on the relative importance of the volume and intensity dimensions of PA,
311 and thus be of importance for future PA research and guideline development. Rowlands et al. [5]
312 aimed to develop the intensity gradient as a metric that compared to traditional summary measures
313 of PA intensity was less dependent on the overall PA level. It has been shown in several studies that
314 associations between overall PA level and the intensity gradient are considerably weaker ($r = 0.36$ –
315 0.56) than between overall PA level and MVPA ($r = 0.93$ – 0.96), which suggest the intensity gradient is
316 more reflective of the intensity *per se* than summary measures of PA intensity [5, 11, 12]. However,
317 we found much stronger associations between overall PA and the intensity gradient in both our

318 datasets ($r = 0.73\text{--}0.86$) than found in previous studies. The use of raw acceleration data in previous
319 studies versus count data used herein likely explains the findings. The frequency dependent filtering
320 used in the generation of ActiGraph counts attenuates capture of high intensity activity reducing
321 associations between the intensity spectrum and cardiometabolic health [34]. This has direct
322 implications for the intensity gradient, which is sensitive to even very small amounts of high intensity
323 activity [35]. Consequently, we observed that the intensity gradient was strongly associated with
324 PC_{Volume} ($r = 0.90\text{--}0.91$), but weakly associated with $PC_{\text{Intensity}}$ ($r = -0.41\text{--}0.40$), which indicates the
325 intensity gradient was not primarily a measure of intensity in the present study. Notably, the
326 collinearity of the intensity gradient and overall PA restricted us from including these variables in
327 joint multiple linear regression models, which may have resulted in poorer model performance than
328 for the principal components analysis for which both volume and intensity components were
329 included.

330 We are not aware of previous studies that have used principal component analysis for investigating
331 the structure of the PA intensity spectrum. The structure of the two datasets included in the present
332 analysis was similar: For PC 1, a higher score indicate a child exhibit more PA and less SED (i.e.,
333 indicative of PA volume), while for PC 2, a higher score means a child have relatively more light
334 intensity PA and relatively less high intensity PA (i.e., indicative of PA intensity). Thus, our findings
335 suggest both higher volume and higher intensity are favourably associated with cardiometabolic
336 health in the ASK dataset, whereas only higher volume was favourably associated with motor skills in
337 the PRESPAS dataset. The latter finding might be counterintuitive given that the strongest association
338 with motor skills were found for 10000–10999 cpm, which could be interpreted as spending time at
339 very high intensities, as opposed to lower intensities, would be favourable to develop motor skills.
340 Notably, it can be observed that high intensities (5000–7999 and 8000–10999 cpm in the ASK and
341 PRESPAS datasets, respectively) have the highest loadings for PC_{Volume} in both datasets, which means
342 these variables contribute most to the overall volume of PA. Although not immediately intuitive, this
343 finding may be reasonable given that time spent at higher intensities will lead to accumulation of

344 much more counts than time spent at lower intensities (e.g., 1 minute spent at 10000 cpm will
345 accumulate as many counts as 100 minutes spent at 100 cpm). Thus, time spent in higher intensities
346 will inherently contribute largely to the volume of PA, as determined by average counts per minute
347 or average acceleration, which is consistent with our findings from the principal component analysis.
348 Thus, despite we extracted two apparently interpretable PCs, the volume and intensity dimensions of
349 PA might still be difficult to separate and apply. This point may also be illustrated by the finding that
350 PC_{Volume} explained 62.8–69.0% of the total variation among the PA variables, whereas $PC_{Intensity}$ only
351 explained 14.4–14.8% of this variation. This finding shows that the relative intensity distribution only
352 constitute a minor part of the overall PA data structure.

353 While the association pattern derived from the multivariate pattern analysis shown for
354 cardiometabolic health in the ASK dataset was similar to the pattern shown previously (using 1-
355 second epoch data) [18], we observed the strongest associations for motor skills in the PRESPAS
356 dataset for 10000–10999 cpm herein compared to 6000–6999 cpm observed previously [14]. Since
357 the intensity gradient is constructed using log-transformed data [5] and since log-transformed (and
358 log-centred) data has been shown to improve model fit compared to raw data [2], all analyses in the
359 present study were based on log-transformed raw data. The variable distributions are typically
360 positively skewed for the highest PA intensities. Skewed data may lead to a problem for modelling
361 since validation and optimization of model selection (i.e., the number of PLS components included) is
362 based on repeated Monte-Carlo resampling. The procedure use half of the sample for modelling and
363 half of the sample for prediction, randomly partitioned for each repetition. Skewed distributions at
364 the higher end of the PA intensity spectrum means that several PLS components that are weakly
365 associated with the predicted outcome are needed to accommodate this variation between
366 participants. The use of log-transformed data makes the distributions for these higher PA intensities
367 less skewed, and thus more stable to resampling, which ultimately leads to simpler and more robust
368 descriptions of data. This effect has probably led to stronger associations for the highest intensities in
369 the PRESPAS dataset, for which we included the most detailed description of the highest intensities

370 (up to ≥ 15000 cpm). This finding could indicate that very high intensity or impact activities, possibly
371 accrued through early sport participation, are the strongest markers of young children's motor
372 development.

373

374 Strengths and limitations

375 The main strength of the present study is the direct comparison of different analytic approaches to
376 analyze associations between PA intensity spectra and two different outcomes in two large datasets.
377 The use of these two datasets allowed for robust comparisons of the statistical approaches, and
378 provided a nuanced picture of the findings beyond what would be possible with only one dataset.
379 Importantly, the structure of the datasets with respect to inter-relationships between variables and
380 extraction of PCs were similar, which illustrates stability and consistency of the findings.

381 The cross-sectional designs limit our ability to draw conclusions about causality. It should also be
382 kept in mind that use of other cohorts, for example spanning other age groups, and the use of other
383 outcomes, could lead to other findings due to different correlation structures among the explanatory
384 PA variables and/or different association patterns between PA intensities and outcomes. The use of
385 waking time count data herein compared to the use of 24-hour raw acceleration data in previous
386 studies [5, 11, 12] could possibly influence the performance of the intensity gradient. Yet, this is the
387 first time the intensity gradient is calculated using waking time count data, which improves our
388 understanding of its features as applied to various types of data. Further studies are warranted to
389 explore these analytic issues and extend our findings.

390

391 Conclusion

392 Our results demonstrate broadly consistent findings are evident across all three analytical
393 approaches. The use of high-resolution PA intensity spectra for determination of associations with
394 outcomes may circumvent limitations imposed by the use of a priori defined intensity cut points and
395 improve the information obtained from accelerometry beyond that of traditional summary measures
396 of intensity. We compared multivariate pattern analysis, which can handle the multicollinearity
397 among variables and thus retain all the information in the data, with dimension reduction methods
398 that can be used to reduce the intensity spectrum to simpler metrics, for determining associations
399 with health and development outcomes in children. Our findings suggest that multivariate pattern
400 analysis explains the most variance in outcomes since it is able to retain information from the data
401 that is lost in other approaches. Yet, the intensity gradient provided the best descriptor of the data
402 using one single metric. Thus, both multivariate pattern analysis and the intensity gradient are
403 preferred over the traditional summary measure approach, depending on the application. Finally, our
404 results suggest volume and intensity dimensions of PA are inherently related and thus not easily
405 disentangled. Principal component analysis might therefore have limited application in association
406 analysis of spectrum PA descriptions.

407

408 Data availability

409 The datasets used in the current study are available from the corresponding author on reasonable
410 request.

411

412 Disclosure of interests

413 The authors declare that they have no competing interests.

414

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420

421 Authors' contributions

422 EAA developed the idea of the study. EAA and AKON collected the data. EAA and OMK designed the
423 study and analyzed the data. EAA wrote the manuscript draft. All authors discussed the
424 interpretation of the results, and read and approved the final manuscript.

425

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434

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538

539 **Figure Legends**

540 **Figure 1. Factor loadings for physical activity intensity variables on the two principal components**
541 **extracted from the principal component analysis.** The total explained variances of the two principal
542 components were 77.3 and 83.8% in the ASK and PRESPAS datasets, respectively.

543 **Figure 2. Association patterns between physical activity intensities and a composite**
544 **cardiometabolic health score (ASK dataset) and locomotor skills (PRESPAS dataset).** Models
545 included 6 and 2 PLS components, respectively. Selectivity ratios are calculated as explained to total
546 variance on the predictive (target projected) component.

547