

1 **Interpretation of associations between the accelerometry physical activity spectrum and**
2 **cardiometabolic health and locomotor skills in two cohorts of children using raw, normalized, log-**
3 **transformed, or compositional data**

4 Eivind Aadland (eivind.aadland@hvl.no),¹ Lars Bo Andersen (lars.bo.andersen@hvl.no),¹ Jairo Hidalgo
5 Migueles (jairoh@ugr.es),² Francisco B Ortega (ortegaf@ugr.es),^{2,3} Olav Martin Kvalheim
6 (olav.kvalheim@uib.no)⁴

7
8 ¹*Western Norway University of Applied Sciences, Faculty of Education, Arts and Sports,*
9 *Department of Sport, Food and Natural Sciences, Campus Sogndal, Sogndal, Norway*

10 ²*PROFITH “PROmoting FITness and Health through physical activity” Research Group, Sport and*
11 *Health University Research Institute (iMUDS), Department of Physical and Sports Education, Faculty*
12 *of Sport Sciences, University of Granada, Granada, Spain*

13 ³*Department of Biosciences and Nutrition, Karolinska Institutet, Huddinge, Sweden.*

14 ⁴*University of Bergen, Department of Chemistry, Bergen, Norway*

15

16

17 **Corresponding author**

18 Eivind Aadland

19 Western Norway University of Applied Sciences, Faculty of Education, Arts and Sports,

20 Department of Sport, Food and Natural Sciences, Campus Sogndal, Box 133, 6851 Sogndal, Norway.

21 Phone: +47 5767 6086; Email: eivind.aadland@hvl.no

22

23

24 **Word count main text: 5044; abstract: 199**

25

26 **Abstract**

27 It is discussed whether associations between accelerometer-derived physical activity intensities and
28 outcomes should be analyzed as absolute or relative data. The aim of the present study was to
29 compare interpretation of association patterns of spectrum physical activity descriptions with
30 outcome using raw, normalized, log-transformed, or compositional data. We used two datasets
31 including 1) 841 schoolchildren and a cardiometabolic health outcome and 2) 1081 preschool
32 children and a locomotor skill outcome. Accelerometry (ActiGraph GT3X+) data were described using
33 multiple variables across the intensity spectrum. We varied the binning of variables to examine
34 sensitivity of the compositional analyses to changes in the distribution center. We used multivariate
35 pattern analysis for all analyses and interpretations of data. Analyses of absolute (i.e., non-
36 compositional) data showed weak associations for lower intensities and strongest associations with
37 cardiometabolic health and locomotor skills for vigorous intensities. The same association patterns
38 were partly observed for the compositional data, but association patterns were in some cases
39 conflicting. The binning of variables had a major influence on associations for compositional data, but
40 not for absolute data, meaning that conclusions depend on the operationalization of compositional
41 data. These differences challenge and confuse interpretation of association patterns derived from
42 the different approaches.

43 **Keywords** Multivariate pattern analysis; Compositional data analysis; Multicollinearity; Singularity,
44 Log-transform, Log-ratio; Children; Accelerometer

45

46 **Background**

47 Accelerometers capture movement on an absolute scale, from which time spent in different
48 intensities is commonly derived. If defining a measurement period as finite, for example the wear
49 period or the 24 hours of a day, movement behaviors can be considered compositional. According to
50 this view, each behavior only contains information relative to the whole composition [1, 2]. Thus,
51 within a fixed period, varying time spent in different activities will substitute each other and be
52 reallocated, for example along the physical activity (PA) intensity spectrum. This feature of “closure”
53 has recently stimulated discussion about the best way to handle and analyze accelerometer-derived
54 PA data described by multiple correlated variables [3]. Both isotemporal substitution models [4] and
55 compositional analysis [1, 2] have been suggested as possible approaches to handle this closure or
56 constant-sum constraint (see Aadland et al [3] for a brief review of the methods). However, both
57 approaches commonly rely on linear least squares regression analysis as their statistical
58 underpinning. Thus, although these approaches theoretically treat PA data as compositional, and
59 thus solve the closure problem with respect to reallocation of time across variables, linear regression
60 cannot handle the much greater challenge of strong multicollinearity resulting from activity patterns
61 inducing correlation patterns among PA variables. This multicollinearity makes linear regression
62 analysis inappropriate for analysis of such data [5, 3], already with few explanatory variables.

63 The deficiency of linear regression is further exaggerated by the use of more detailed PA descriptions
64 as compared to traditional blunt descriptions typically using 3–4 variables [3], for example sedentary
65 time (SED), light PA (LPA), moderate-to-vigorous PA (MVPA), and/or sleep. In line with recent
66 suggestions for an improved use of accelerometer data [6-8], Aadland et al recently showed that the
67 inclusion of the entire intensity spectrum and using an improved resolution of the PA description
68 greatly increased information derived from accelerometry, and thus knowledge about associations
69 between PA and cardiometabolic health, in children [9, 10]. These findings show the unexploited
70 potential of accelerometry when moving from summary measures to more detailed descriptions of

71 PA. Yet, such data requires statistical methods that are able to handle this type of highly correlated
72 data. Therefore, Aadland et al introduced multivariate pattern analysis to analyze associations
73 between the multicollinear explanatory PA variables and the cardiometabolic health outcomes [9,
74 10]. Multivariate pattern analysis is widely applied in other fields of research with the objective of
75 revealing patterns of important biomarkers among hundreds or thousands of highly interrelated
76 variables [11-13], and can handle completely collinear explanatory variables using latent variable
77 modelling [14, 15]. Since multivariate pattern analysis can handle multicollinear data, Aadland et al
78 [3] were able to circumvent the limitations of linear regression and directly compare association
79 patterns between raw (min/day) and compositional PA data with cardiometabolic health. Both a
80 traditional description reducing the PA profiles to four variables (SED, LPA, moderate PA (MPA), and
81 vigorous PA (VPA)) and a spectrum description of 23 variables were used. With both descriptions we
82 found apparently differing association patterns with cardiometabolic health using raw or
83 compositional data. We also found that models using compositional data lead to higher explained
84 variances than models using raw data, which could be attributed to reduction of noise due to log-
85 transformation [16]. However, it was not examined whether the improved model fit resulted from
86 the log-transformation or from the centering of data. Thus, it should be examined whether the
87 improved model fit is a favorable feature of compositional data compared to raw and/or log-
88 transformed data.

89 We did not provide a comprehensive elaboration on interpretation of the findings from raw or
90 compositional data in the previous paper [3], which is crucial to fully understand the results and thus
91 inform PA guidelines and provide direction for future research. Thus, a broader exploration and
92 discussion of the interpretation of the absolute (raw, normalized, and log-transformed data) and
93 relative (compositional data) nature of these association patterns are needed. The aim of this paper
94 is therefore to extend our previous analysis [3] on comparison of results and interpretation of
95 associations using raw, normalized, log-transformed, and compositional PA data. For improved

96 generalization, we will analyze two large datasets (in preschool- and schoolchildren) and two
97 different outcomes (cardiometabolic health and locomotor skills) in this work.

98

99 **Methods**

100 We have previously published the PA signature associated with cardiometabolic health in the *Active*
101 *Smarter Kids (ASK)* study [9, 10, 3] and the PA signature associated with locomotor skills in *The Sogn*
102 *og Fjordane Preschool Physical Activity Study (PRESPAS)* [17]. The aim of the present study is limited
103 to *compare* association patterns using raw and compositional data within these datasets. We refer
104 readers to previously published descriptions of sampling and children’s characteristics, study
105 protocols, instruments, and procedures of the ASK study [9, 10, 3, 18] and the PRESPAS study [19, 17]
106 for detailed study information. Thus, we provide below only a brief overview of the most relevant
107 information to provide sufficient context to support the study aim of comparing association patterns
108 between models relying on different treatment of the explanatory data matrix.

109

110 **Participants**

111 The ASK study was conducted in western Norway during 2014–2015 and included 841 10-year old
112 schoolchildren providing relevant explanatory (PA) and outcome (cardiometabolic health) data [9, 10,
113 3, 18]. The PRESPAS study was conducted in western Norway during 2015–2016 and included 1081 3-
114 6-year old preschool children providing relevant explanatory (PA) and outcome (locomotor skills)
115 data [19]. Procedures and methods in both studies conform to ethical guidelines defined by the
116 World Medical Association’s Declaration of Helsinki and its subsequent revisions. . The Norwegian
117 South-East Regional Committee for Medical Research Ethics and the Norwegian Centre for Research
118 Data approved the study protocols. We obtained written informed consent from each child’s parents
119 or legal guardians and from the responsible preschool and school authorities prior to all testing.

120
121
122
123
124
125
126
127
128
129
130
131
132
133
134
135
136
137
138
139
140
141
142
143
144

Procedures

Physical activity

PA was measured using the ActiGraph GT3X+ accelerometer (Pensacola, FL, USA) [20] worn at the waist over seven (ASK) and 14 (PRESPAS) consecutive days, except during water activities (swimming, showering) or while sleeping. Units were initialized at a sampling rate of 30 Hz and files were analyzed restricted to hours 06:00 to 23:59 using 1-second epochs to capture low and high intensity PA [21] using the KineSoft analytical software version 3.3.80 (KineSoft, Loughborough, UK). We applied wear time requirements of ≥ 8 hours/day and ≥ 4 days/week to constitute a valid measurement [22, 23].

As described previously, we applied “spectrum descriptions” of PA to obtain a better and more nuanced picture of associations between PA and the outcomes compared to traditional blunt descriptions of data. We have previously used descriptions of 23 variables (from 0–99, 100–249, 500–999, 1000–1499, ... 9500–9999, to ≥ 10000 cpm) in the ASK dataset [3] and 33 variables (from 0–99, 100–249, 500–999, 1000–1499, ... 14500–14999, to ≥ 15000 cpm) in the PRESPAS dataset [17], obtained from the vertical axis, to capture movement in narrow intensity intervals across the intensity spectrum. Thus, except for the two lowest bins and the highest bin, intervals of 500 cpm were used in the original studies. For simplicity, we created new bins with intervals of 1000 cpm for both datasets herein. Thus, the ASK data was described using 12 bins (from 0–99, 100–999, 1000–1999, 2000–2999, ... 9000–9999, to ≥ 10000 cpm) and the PRESPAS data was described using 17 bins (from 0–99, 100–999, 1000–1999, 2000–2999, ... 14000–14999, to ≥ 15000 cpm). These descriptions performed similarly to previously published models using higher resolution (explained variances = 17.02% for 500 cpm intervals and 15.69% for 1000 cpm intervals in the ASK dataset and 6.70% for 500 cpm intervals and 6.84% for 1000 cpm intervals in the PRESPAS dataset). In addition, we used the original descriptions of 23 and 33 bins in the ASK and PRESPAS datasets, respectively, as a basis

145 for examining the sensitivity of the analyses to different resolution of the data across the intensity
146 spectrum. Specifically, in each dataset, we constructed descriptions that were condensed to 1) three
147 bins *below* 4000 cpm (0–99, 100–1999, and 2000–3999), where the original bins above 4000 cpm
148 were retained, and 2) three bins *above* 4000 cpm (4000–5999, 6000–7999, and ≥ 8000 cpm in the
149 ASK dataset; 4000–7999, 8000–11999, and ≥ 12000 cpm in the PRESPAS dataset), where the original
150 bins below 4000 cpm were retained.

151 We used the Evenson et al [24, 25] intensity cut points of 0–99, 100–2295, 2296–4011, and ≥ 4012
152 cpm as a guidance for interpreting intensities across the spectrum as SED, LPA, MPA, and VPA *post*
153 *hoc*.

154

155 Outcomes

156 The outcome in the ASK study was a cardiometabolic composite score [26] calculated as the mean of
157 six variables (systolic blood pressure, triglycerides, total:high-density lipoprotein cholesterol,
158 homeostasis model assessment of insulin resistance, waist:height ratio, and the inverse Andersen
159 aerobic fitness test) using standardized scores after adjustment for sex and age using residuals from
160 linear regression. A higher score indicates poorer cardiometabolic health.

161 The outcome in the PRESPAS study was a sum score of three locomotor movement tasks (run,
162 horizontal jump, hop) guided by the Test of Gross Motor Development 3 test battery [27, 28]. A
163 higher score indicates better locomotor skills. Children were scored quantitatively based on whether
164 they did or did not demonstrate specific criteria for each skill based on the original scoring
165 procedures. The criteria scores were averaged for each task and the total locomotor score. The score
166 was standardized after adjustment for sex, age, body mass index, preschool, and assessor of motor
167 skills using residuals from linear regression prior to analysis.

168

169 Statistical analyses

170 We analyzed PA profiles as raw data (i.e., min/day), normalized data (proportion of wear time), log-
171 transformed data, and as compositional data. Compositional transformation was performed using
172 the centered log-ratio (clr) method as described by Hinkle and Rayens [29] and by constructing log-
173 ratios according to SED (i.e., 0–99 cpm, SED log-ratio (SEDIr)), LPA (i.e., 1000–1999 cpm, LPAIr), MPA
174 (i.e., 3000–3999 cpm, MPAIr), and VPA (i.e., the variable strongest associated with the outcome using
175 absolute data = 7000–7999 cpm in the ASK dataset and 5000–5999 cpm in the PRESPAS dataset,
176 VPAIr). Because multivariate pattern analysis can handle any degree of multicollinearity, the use of
177 the isometric log-ratio [30] was not necessary (the isometric log-ratio technically provides an open
178 dataset and makes it possible to analyze data by linear regression in the absence of other sources of
179 multicollinearity than closure, i.e., a correlation of -1 spread over the explanatory variables).
180 Moreover, the isometric log-ratio transform does not allow for determination of the association
181 pattern from one joint interpretable model (i.e., separate models has to be performed for each
182 intensity bin). Centred log-ratios were made by 1) normalization of the PA intensity profile for each
183 individual, that is, calculation of proportions where all explanatory variables sum to 1, prior to 2)
184 making natural log-transformations of each variable, and 3) centering the explanatory variables for
185 each object to the mean logarithm of all explanatory variables (clr) [31]. Construction of other log-
186 ratios (SEDIr, LPAIr, MPAIr, VPAIr) followed steps 1 and 2, but used the selected PA bins (SED, LPA,
187 MPA, and VPA) as the denominator.

188 *Multivariate pattern analysis.* Partial least squares (PLS) regression analysis [14] was used to
189 determine the multivariate association patterns of raw and compositional PA data (explanatory
190 variables) with cardiometabolic health (ASK dataset) and locomotor skills (PRESPAS dataset)
191 (outcomes) (please see Aadland et al [3, 32] for a brief overview of this statistical approach as applied
192 to PA accelerometry data). PLS regression decomposes the explanatory variables into orthogonal
193 linear combinations (PLS components), while simultaneously maximizing the covariance with the

194 outcome variable. Thus, PLS regression is able to handle completely collinear variables through the
195 use of latent variable modelling [14]. The procedure differs from that of factor analysis or principal
196 component analysis by creating components that maximize the covariation with the outcome, not
197 internally among the explanatory variables. Prior to PLS regression, all variables were centered and
198 standardized to unit variance. Models were cross-validated using Monte Carlo resampling with 1000
199 repetitions by repeatedly and randomly keeping 50% of the subjects as an external validation set
200 when estimating the models to determine the number of PLS components [33]. Validation is an
201 integrated part of the procedure to avoid overfitting due to inclusion of minor PLS components
202 representing noise. For each validated PLS regression model, a single predictive component was
203 subsequently calculated by means of target projection [15, 11] to express all the predictive variance
204 in the PA intensity spectrum related to cardiometabolic health in a single intensity vector. Selectivity
205 ratios (SRs) with 95% CIs were obtained as the ratio of this explained predictive variance to the total
206 variance for each PA intensity variable [32, 34, 35]. Thus, the SRs provide the explained variance of
207 each variable with the predicted outcome in the multivariate space, while retaining their direction of
208 association. The procedure for obtaining the multivariate patterns is completely data-driven, with no
209 assumptions on variable distributions or degree of collinearity among variables. These analyses were
210 performed by means of the commercial software Sirius version 11.0 (Pattern Recognition Systems
211 AS, Bergen, Norway).

212 Geometric means were calculated for the mean logarithm of all explanatory variables to compare the
213 center of different clr models. Pearson's correlation was used to compare association patterns for
214 raw, normalized, and log-transformed data.

215

216 **Results**

217 We included 841 schoolchildren (mean (SD) 10.2 (0.3) years old, 50% boys) and 1081 preschool
218 children (4.7 (0.9) years old, 52% boys) who provided valid data on all relevant variables.

219 Figures 1 and 2 show the multivariate association patterns between PA and cardiometabolic health
220 (ASK dataset) and between PA and locomotor skills (PRESPAS dataset), respectively, using raw
221 (min/day), normalized (proportion of wear time), log-transformed, and compositional data. Within
222 both datasets, explained variances were similar for raw and normalized data, but improved for log-
223 transformed and compositional data. Still, association patterns were similar for models using
224 absolute (i.e., raw, normalized, and log-transformed) data (ASK: $r \geq 0.99$; PRESPAS: $r \geq 0.93$). In the
225 ASK dataset, the strongest association with cardiometabolic health was found for 7000–7999 cpm
226 when using absolute data. Consistent with this finding, the compositional data also showed
227 reallocating time from lower intensities to 7000–7999 cpm was most favorable. In contrast, while
228 5000–5999 cpm was strongest associated with locomotor skills in the PRESPAS dataset when using
229 absolute data, compositional data showed reallocating time from lower intensities to 12000–12999
230 cpm was most favorable. The centers (geometric means) of the compositional data were 0.0163 and
231 0.0070 in the ASK and PRESPAS datasets (Supplemental Figure 1), respectively, having corresponding
232 turning points for favorable/unfavorable reallocation of time around 4000–4999 and 7000–7999 cpm
233 (Figures 1 and 2).

234 To further explore the possible impact of including higher intensities and having a higher number of
235 variables in the PRESPAS dataset (0–99 to ≥ 15000 cpm; 17 variables) than in the ASK dataset (0–99
236 to ≥ 10000 cpm; 12 variables), we tested association patterns using datasets with different resolution
237 of data across the intensity spectrum. In both the ASK dataset (Figure 3) and the PRESPAS dataset
238 (Figure 4), varying the resolution had a clear impact on geometric means and association patterns
239 using compositional data. In the ASK dataset, the turning points (for which time should be
240 reallocated around) were 5500–5999 (geometric mean = 0.0060) and 3000–3499 cpm (geometric
241 mean = 0.0249) when using datasets that were condensed below and above 4000 cpm, respectively.
242 This difference was more pronounced in the PRESPAS dataset, for which corresponding turning
243 points of 9500–9999 (geometric mean = 0.0023) and 2500–2999 cpm (geometric mean = 0.0270)
244 were found.

245 In addition to using the data-driven centered log-ratio (which use the geometric mean composition
246 as its reference for constructing log-ratios), we explored how the use of alternative log-ratios (SEDIr,
247 LPAI_r, MPAI_r, and VPAI_r) performed. In the ASK dataset (Figure 5), reallocating time from SED (0–99
248 cpm) and LPA (2000–2999 cpm) to all other intensities were favorable, whereas the findings for MPA
249 (3000–3999 cpm) were mixed (i.e., favorable to reallocate time from lower intensities to MPA and
250 unfavorable to reallocate time from higher intensities to MPA). Reallocating time from VPA (i.e., the
251 intensity strongest associated with cardiometabolic health using absolute data; 7000–7999 cpm) to
252 all other intensities were unfavorable. In the PRESPAS dataset (Figure 6), the association patterns
253 were rather similar to those found in the ASK dataset. However, when using VPAI_r (i.e., the intensity
254 strongest associated with locomotor skills using absolute data (5000–5999 cpm) as the denominator
255 for constructing compositions), the results showed reallocating time from 5000–5999 cpm to higher
256 intensities was favorable, and thus similar to findings for clr, but in conflict with the interpretation
257 using absolute data.

258

259 **Discussion**

260 In the present study we used two large datasets including children of different age to explore
261 association patterns between two different outcomes and spectrum descriptions of PA using raw,
262 normalized, log-transformed, and compositional data. Our findings provide a good basis for
263 elaboration on interpretation of different models, and show some similarities, but also highlight
264 some challenges and dissimilar interpretations. Briefly, while all findings show that intensities in the
265 vigorous range is strongest associated with the outcomes in both datasets, the relative nature of the
266 compositional data leads to some counterintuitive findings and differences in interpretation caused
267 by the operationalisation of PA data. These findings challenge and confuse interpretation of
268 association patterns derived from the different approaches. Our findings therefore suggest future

269 studies using accelerometry-derived PA data apply multivariate pattern analysis with raw data, or
270 possibly log-transformed data, to avoid confusion in developing the field of PA epidemiology.

271 While associations for raw, normalized, and log-transformed data are *absolute* (i.e., associations are
272 given as each variable's importance for the predicted outcome in a multivariate space), associations
273 for compositional data are *relative* (i.e., associations for all variables are relative to other variables).
274 Thus, although the association patterns in Figures 1 and 2 appear different, the interpretation of the
275 patterns are partly equivalent. For the ASK dataset (Figure 1), the use of absolute data shows
276 vigorous intensities are strongly associated with cardiometabolic health, whereas associations for
277 lower intensities are weak. For comparison, the compositional data shows that reallocating time
278 from lower intensities to vigorous intensity is favourable. This interpretation can also be made from
279 the absolute data; since spending more time in VPA is favourable, spending less time in VPA – and
280 thus more time in lower intensities – is unfavourable. However, the results from the compositional
281 data presents three major challenges for interpretation.

282 First, since associations are relative, it is not possible to determine how each variable is associated
283 with the outcome. This challenge also, to some extent, applies to interpretation of isotemporal
284 substitution models. For example, if it is favourable to reallocate time from SED to VPA, it will by
285 definition also be unfavourable to reallocate time from VPA to SED, but does that mean spending
286 more time in VPA is favourable or more time in SED is unfavourable (or both)? The compositional
287 data in Figure 1 clearly suggests spending time in SED, LPA, and MPA are as detrimental to
288 cardiometabolic health as VPA is favourable, because the associations are mirrored around a turning
289 point, that is, the centre of the distribution (the geometric mean composition). In essence, this
290 approach reduces findings to the trivial interpretation that one should substitute all PA activities with
291 an intensity below the centre point to activities with an intensity above this point, because it is not
292 possible to determine the intensities' absolute relation to the outcome. This point is also
293 underpinned by constructing alternative log-ratios (Figures 5 and 6), of which nearly all provide the

294 hypothesized association patterns. For example, while the SEDlr shows reallocating time from SED to
295 higher intensities is favourable, the VPAIr shows reallocating time from VPA to lower intensities is
296 unfavourable. Still, similar to the clr, the VPAIr would easily be misinterpreted and taken as evidence
297 of an unfavourable association between low intensities (up to 6999 cpm) and cardiometabolic health,
298 though this association pattern is relative and trivial.

299 Second, all findings using compositional data suggest it is more unfavourable to substitute time in
300 higher intensities with time in LPA than in SED. This finding contradict current evidence indicating a
301 possible favourable association between LPA and cardiometabolic health [6] but no association for
302 SED and cardiometabolic health [36], and our raw data showing a weak unfavourable association for
303 SED, but no association for LPA. We simply suggest this finding is a statistical artefact resulting from
304 the great alteration of the correlation structure of the explanatory data matrix [3]. This enforced
305 correlation structure might also result in the major shift in the intensities that are most favourably
306 associated with locomotor skills in the PRESPAS dataset (Figure 2). While the strongest association is
307 found for 5000–5999 cpm using absolute data, the analysis of compositional data suggest it is most
308 favourable to spend time in 12000–12999 cpm. We suggest these differing findings result from the
309 nature of the compositional transformation; since associations are relative, associations are not
310 favourable or unfavourable, but rather more favourable or less favourable, which leads to a biphasic
311 association pattern around the geometric mean composition. This turning point will inherently be in
312 the middle of the distribution, which is where we (accidentally) find the strongest absolute
313 association in the PRESPAS dataset. A similar pattern is seen for the VPAIr (Figure 6); since the
314 variable showing the strongest association with locomotor skills was used for constructing the log-
315 ratio, we would expect reallocating time from this intensity to all others intensities would be
316 unfavourable. In contrast, reallocating time to higher intensities seems favourable. These findings
317 result from the enforced biphasic association pattern, driven by the nature of the log-ratio.
318 Interestingly, the geometric mean composition essentially provide the same information as the
319 intensity gradient, introduced by Rowlands et al [37], as both measures capture and condense the

320 distribution of time spent across the intensity spectrum to a single variable (r between intensity
321 gradient and geometric mean = 0.96 in the ASK dataset and 0.97 in the PRESPAS dataset, results not
322 reported). A higher geometric mean means more time is spent in higher intensities, which means the
323 slope across the intensity spectrum is less steep. Thus, a higher geometric mean means the intensity
324 distribution is moved upwards, resulting in lower intensities being negatively related to the mean
325 and higher intensities being positively related to the mean using compositional data. This biphasic
326 pattern, suggesting it is favourable to reallocate time from lower to higher intensities, is also seen
327 when restricting the spectrum in PRESPAS to intensities above 5000 cpm (results not shown), for
328 which weaker associations are seen for increasing intensities using absolute data (Figure 2). Thus,
329 these approaches result in directly conflicting findings.

330 Third, another challenge for interpretation of compositional data is the major influence on findings
331 resulting from different operationalisation or binning of PA data. As shown in Figures 3 and 4,
332 different binning – of the same data – result in largely different interpretations. We hypothesized the
333 clr would be vulnerable to the operationalisation of data, because the number of variables and their
334 compositions would affect the geometric mean and thus the centre of the distribution. While the
335 influence was moderate in the ASK dataset (the centre was found for 3000–3499 cpm versus 5500–
336 5999 cpm for the datasets condensed above and below 4000 cpm, respectively), a major influence
337 was found in the PRESPAS dataset (the centre was found for 2500–2999 cpm versus 9500–9999 cpm
338 for the datasets condensed above and below 4000 cpm, respectively) for which a higher number of
339 higher intensity variables was included. Thus, more or less arbitrary choices in variable selection or
340 binning may lead to substantial differences in interpretation, which may confuse conclusions and
341 hamper comparability between studies.

342 An assumption underlying the possible need for a compositional transformation of data, which, to
343 the best of our knowledge, has not been discussed previously, is how PA variables across the
344 intensity spectrum are associated with the outcome. The general idea behind constructing a closed

345 dataset is that time needs to be reallocated among activities or intensities, that is, time cannot be
346 added to some intensities without detracting from other intensities. However, most of the time over
347 a day is spent sedentary or in low intensity, which is not important to health and developmental
348 outcomes in children [9, 3, 17, 36]. Thus, there is arguably abundant time that that can be added to
349 the higher intensities without influencing the association model. In the included datasets, 719 out of
350 795 minutes/day (ASK) and 627 out of 702 minutes/day (PRESPAS) (89–90% of the time) is
351 accumulated in SED and LPA [3, 17], for which associations with the outcomes are very weak. Thus, it
352 does not matter if some minutes are taken from the lower intensities to increase time in higher
353 intensities. This point challenges the foundational assumption of compositional analysis and support
354 the use of absolute data for analysis.

355 We have previously shown a great improvement of association models between PA and
356 cardiometabolic health when using a more detailed PA description compared to traditional
357 descriptions [9, 10]. This improved resolution and higher number of variables have several important
358 implications for analysis. First, if using a blunt description of only two variables, for example SED and
359 non-SED PA, given a constant sum of these variables (i.e., total wear time), they will be perfectly
360 negatively correlated and thus singular (the imposed correlation of the explanatory data matrix is -1).
361 In this situation, spending more time in one variable will lead to an equivalent reduction in time
362 spent in the other, so time will be fully reallocated across these variables. In situations with few
363 variables, this reallocation might be a concern. However, when analyzing an explanatory data matrix
364 comprising a higher number of variables, reallocation of time will be spread over many variables and
365 thus be less important when estimating associations [3, 32]. This feature makes the transformation
366 to compositional data less relevant in this situation than for traditional descriptions of data (for
367 example SED, LPA, and MVPA, or different types of movement). Also with few variables, though,
368 whether raw or compositional, data will be correlated, which leaves linear regression less suited for
369 analysis [3, 5]. Importantly, if using multivariate pattern analysis, the construction of compositions is
370 not needed, since the model can handle singular data. Interestingly, though, log-transformation of

371 data (centered or not) lead to improved model fit in both datasets, as also demonstrated for
372 compositional data using both a traditional and spectrum description previously [3]. This finding
373 suggests log-transformation reduces noise in the accelerometer data. It is well-known that count
374 data produces heteroscedastic noise, which means that noise increases with the number of counts
375 [16]. This source of noise is largest for SED and LPA. At the other end of the intensity spectrum, the
376 number of counts is much lower, but the distributions are typically positively skewed. Skewed data
377 may lead to a problem for modelling since validation and optimization of model selection (i.e., the
378 number of PLS components included) is based on repeated Monte-Carlo resampling. The procedure
379 use half of the sample for modelling and half of the sample for prediction, randomly partitioned for
380 each repetition. Skewed distributions at the higher end of the PA intensity spectrum means that
381 several PLS components that are weakly associated with the predicted outcome are needed to
382 accommodate this variation between participants. The use of log-transformed data makes the
383 distributions for these higher PA intensities less skewed, and thus more stable to resampling, which
384 ultimately leads to simpler and more robust descriptions of data. This effect was most evident in the
385 PRESPAS dataset for which we used the most detailed description of the highest intensities (up to \geq
386 15000 cpm) (Figure 2).

387

388 Strengths and limitations

389 The main strength of the present study is the direct comparison of different analytic approaches to
390 analyze associations between PA and two different outcomes in two large datasets. The use of these
391 different datasets allowed for robust comparisons across the statistical approaches, and provided a
392 nuanced picture of the findings beyond what would be possible with only one dataset. Importantly,
393 we included two spectrum descriptions of PA, which, compared to traditional blunt descriptions,
394 provide a better opportunity to reveal how the different handling of data affect the interpretation.
395 Importantly, the same challenges as revealed herein apply to fewer variables, although less apparent.

396 Nevertheless, while a spectrum description may provide a substantially improved picture of
397 association patterns than traditional description with fewer variables [9, 10], a simpler description
398 might be needed for translation of findings into PA guidelines [32].

399 The cross-sectional designs limits our ability to draw conclusions about causality. It should also be
400 kept in mind that use of other cohorts, for example spanning other age groups, and the use of other
401 outcomes, could lead to other findings due to different correlation structures among the explanatory
402 PA variables and/or different association patterns between PA intensities and outcomes. Further
403 studies are therefore warranted to explore these analytic issues and extend our findings. Finally, we
404 do not know the “true” association pattern between PA intensities and cardiometabolic health and
405 motor skills. Thus, our conclusions of which approach provide the best results need to be interpreted
406 with this issue in mind.

407

408 Conclusion

409 Interpretation of associations between accelerometer-derived PA spectra and cardiometabolic health
410 and motor skills in two different samples of children differed when analyzing absolute (raw,
411 normalized, and log-transformed) and relative (compositional) data. While we find interpretation of
412 association patterns for absolute data meaningful, we find interpretation of association patterns for
413 compositional data challenging and partly in conflict with results from absolute data. Moreover, our
414 findings show that the relative nature of compositional data makes interpretation of association
415 patterns susceptible to change according to how the explanatory data matrix is operationalized.
416 Consistent with a previous study [3], we therefore recommend future studies using accelerometry-
417 derived PA data apply multivariate pattern analysis with raw data, or possibly log-transformed data,
418 to avoid confusion in developing the field of PA epidemiology. Finally, we find the use of absolute
419 data more meaningful for development and messaging of PA guidelines, as results can be interpreted
420 directly according to the strength of association between the PA variables and the outcome, and

421 directly in minutes of PA spent in given intensities of activity, although this is complicated by the
422 correlated data structure [32].

423

424 **Funding**

425 The ASK study was funded by the Research Council of Norway (grant number 221047/F40) and the
426 Gjensidige Foundation (grant number 1042294). The PRESPAS study was funded by the Sogn og
427 Fjordane County Municipality. Jairo H. Migueles is supported by a Grant from the Spanish Ministry of
428 Education, Culture and Sport (FPU15/02645). Additional funding was obtained from the
429 SmarterMove project supported by the MINECO/FEDER (DEP2016-79512-R) and the University of
430 Granada, Plan Propio de Investigación 2016, Excellence actions: Units of Excellence; Scientific
431 Excellence Unit on Exercise and Health (UCEES). Junta de Andalucía, Consejería de Conocimiento,
432 Investigación y Universidades and European Regional Development Funds (ref. SOMM17/6107/UGR).
433 None of the funding agencies had any role in the study design, data collection, analyzing or
434 interpreting data, or in writing the manuscripts.

435

436 **Acknowledgements**

437 We thank all children, parents and staff at the participating preschools (PRESPAS) and schools (ASK)
438 for their excellent cooperation during the data collection. We also thank colleagues and students at
439 the *Western Norway University of Applied Sciences* (formerly *Sogn og Fjordane University College*) for
440 their contribution to the ASK and PRESPAS studies. We thank participants at the *International*
441 *Workshop: A focus on statistical methods to analyse accelerometer-measured physical activity*,
442 Granada, Spain, October 21.-22. 2019 for their valuable perspectives on best practices for analyzing
443 physical activity data.

444

445 **Availability of data and materials**

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

448

449 **Competing interests**

450 The authors declare that they have no competing interests.

451

452 **References**

- 453 1. Chastin SFM, Palarea-Albaladejo J, Dontje ML, Skelton DA. Combined Effects of Time Spent in
454 Physical Activity, Sedentary Behaviors and Sleep on Obesity and Cardio-Metabolic Health Markers: A
455 Novel Compositional Data Analysis Approach. *Plos One*. 2015;10(10).
456 doi:10.1371/journal.pone.0139984.
- 457 2. Dumuid D, Stanford TE, Martin-Fernandez JA, Pedisic Z, Maher CA, Lewis LK et al. Compositional
458 data analysis for physical activity, sedentary time and sleep research. *Stat Methods Med Res*.
459 2018;27(12):3726-38. doi:10.1177/0962280217710835.
- 460 3. Aadland E, Kvalheim OM, Anderssen SA, Resaland GK, Andersen LB. Multicollinear physical activity
461 accelerometry data and associations to cardiometabolic health: challenges, pitfalls, and potential
462 solutions. *Int J Behav Nutr Phys Act*. 2019;16(1). doi:10.1186/s12966-019-0836-z.
- 463 4. Mekary RA, Willett WC, Hu FB, Ding EL. Isotemporal Substitution Paradigm for Physical Activity
464 Epidemiology and Weight Change. *Am J Epidemiol*. 2009;170(4):519-27. doi:10.1093/aje/kwp163.
- 465 5. Cohen J, Cohen P, West SG, Aiken LS. Applied multiple regression/correlation analysis for the
466 behavioral sciences. 3 ed. New York: Routledge; 2003.
- 467 6. Poitras VJ, Gray CE, Borghese MM, Carson V, Chaput JP, Janssen I et al. Systematic review of the
468 relationships between objectively measured physical activity and health indicators in school-aged
469 children and youth. *Appl Physiol Nutr Metabol*. 2016;41(6):S197-S239. doi:10.1139/apnm-2015-0663.
- 470 7. van der Ploeg HP, Hillsdon M. Is sedentary behaviour just physical inactivity by another name? *Int J*
471 *Behav Nutr Phys Act*. 2017;14:8. doi:10.1186/s12966-017-0601-0.

- 472 8. Pedisic Z. Measurement issues and poor adjustments for physical activity and sleep undermine
473 sedentary behaviour research - the focus should shift to the balance between sleep, sedentary
474 behaviour, standing and activity. *Kinesiology*. 2014;46(1):135-46.
- 475 9. Aadland E, Kvalheim OM, Anderssen SA, Resaland GK, Andersen LB. The multivariate physical
476 activity signature associated with metabolic health in children. *Int J Behav Nutr Phys Act*.
477 2018;15(77). doi:10.1186/s12966-018-0707-z.
- 478 10. Aadland E, Kvalheim OM, Anderssen SA, Resaland GK, Andersen LB. The Triaxial Physical Activity
479 Signature Associated with Metabolic Health in Children. *Med Sci Sports Exerc*. 2019;51(10):2173-9.
480 doi:10.1249/mss.0000000000002021.
- 481 11. Rajalahti T, Kvalheim OM. Multivariate data analysis in pharmaceuticals: A tutorial review. *Int J*
482 *Pharm*. 2011;417(1-2):280-90. doi:10.1016/j.ijpharm.2011.02.019.
- 483 12. Rajalahti T, Kroksveen AC, Arneberg R, Berven FS, Vedeler CA, Myhr K-M et al. A multivariate
484 approach to reveal biomarker signatures for disease classification: application to mass spectral
485 profiles of cerebrospinal fluid from patients with multiple sclerosis. *J Proteome Res*. 2010;9(7):3608-
486 20. doi:10.1021/pr100142m.
- 487 13. Madsen R, Lundstedt T, Trygg J. Chemometrics in metabolomics-A review in human disease
488 diagnosis. *Anal Chim Acta*. 2010;659(1-2):23-33. doi:10.1016/j.aca.2009.11.042.
- 489 14. Wold S, Ruhe A, Wold H, Dunn WJ. The collinearity problem in linear-regression - the partial least-
490 squares (pls) approach to generalized inverses. *SIAM J Sci Comput*. 1984;5(3):735-43.
491 doi:10.1137/0905052.
- 492 15. Kvalheim OM, Karstang TV. Interpretation of latent-variable regression-models. *Chemometr Intell*
493 *Lab Syst*. 1989;7(1-2):39-51. doi:10.1016/0169-7439(89)80110-8.
- 494 16. Kvalheim OM, Brakstad F, Liang Y. Preprocessing of analytical profiles in the presence of
495 homoscedastic or heteroscedastic noise. *Anal Chem*. 1994;66(43-51).
- 496 17. Nilsen AKO, Anderssen SA, Loftesnes JM, Johannessen K, Ylvisaaker E, Aadland E. The multivariate
497 physical activity signature associated with fundamental motor skills in preschoolers. *J Sports Sci*.
498 2019;1-9. doi:10.1080/02640414.2019.1694128.
- 499 18. Resaland GK, Moe VF, Aadland E, Steene-Johannessen J, Glosvik Ø, Andersen JR et al. Active
500 Smarter Kids (ASK): Rationale and design of a cluster-randomized controlled trial investigating the
501 effects of daily physical activity on children's academic performance and risk factors for non-
502 communicable diseases. *BMC Public Health*. 2015;15:709-. doi:10.1186/s12889-015-2049-y.
- 503 19. Nilsen AKO, Anderssen SA, Resaland GK, Johannessen K, Ylvisaaker E, Aadland E. Boys, older
504 children, and highly active children benefit most from the preschool arena regarding moderate-to-
505 vigorous physical activity: A cross-sectional study of Norwegian preschoolers. *Prev Med Reports*.
506 2019;14:100837-. doi:10.1016/j.pmedr.2019.100837.

507 20. John D, Freedson P. ActiGraph and Actical physical activity monitors: a peek under the hood. *Med*
508 *Sci Sports Exerc.* 2012;44(1 Suppl 1):S86-S9.

509 21. Aadland E, Andersen LB, Anderssen SA, Resaland GK, Kvalheim OM. Accelerometer epoch setting
510 is decisive for associations between physical activity and metabolic health in children. *J Sports Sci.*
511 2019;1-8. doi:10.1080/02640414.2019.1693320.

512 22. Aadland E, Andersen LB, Skrede T, Ekelund U, Anderssen SA, Resaland GK. Reproducibility of
513 objectively measured physical activity and sedentary time over two seasons in children; Comparing a
514 day-by-day and a week-by-week approach. *Plos One.* 2017;12(12).
515 doi:10.1371/journal.pone.0189304.

516 23. Aadland E, Johannessen K. Agreement of objectively measured physical activity and sedentary
517 time in preschool children. *Prev Med Reports.* 2015;2:635-9.

518 24. Evenson KR, Catellier DJ, Gill K, Ondrak KS, McMurray RG. Calibration of two objective measures
519 of physical activity for children. *J Sports Sci.* 2008;26(14):1557-65. doi:10.1080/02640410802334196.

520 25. Trost SG, Loprinzi PD, Moore R, Pfeiffer KA. Comparison of Accelerometer Cut Points for
521 Predicting Activity Intensity in Youth. *Med Sci Sports Exerc.* 2011;43(7):1360-8.
522 doi:10.1249/MSS.0b013e318206476e.

523 26. Andersen LB, Harro M, Sardinha LB, Froberg K, Ekelund U, Brage S et al. Physical activity and
524 clustered cardiovascular risk in children: a cross-sectional study (The European Youth Heart Study).
525 *Lancet.* 2006;368(9532):299-304. doi:10.1016/S0140-6736(06)69075-2.

526 27. Ulrich D. Test of Gross Motor Development – Third Edition. Ann Arbor, MI: Center on Physical
527 Activity and Health in Pediatric Disabilities, 2013.

528 28. Ulrich DA. Test of Gross Motor Development - Third edition. Examiner's Manual. Austin, Texas:
529 Pro.ed.; 2019.

530 29. Hinkle J, Rayens W. Partial least squares and compositional data: problems and alternatives.
531 *Chemometr Intell Lab Syst.* 1995;30(1):159-72. doi:https://doi.org/10.1016/0169-7439(95)00062-3.

532 30. Hron K, Filzmoser P, Thompson K. Linear regression with compositional explanatory variables. *J*
533 *Appl Stat.* 2012;39(5):1115-28. doi:10.1080/02664763.2011.644268.

534 31. Aitchison J. The statistical analysis of compositional data. *Journal of the Royal Statistical Society,*.
535 1982;44(2):139-77.

536 32. Aadland E, Andersen LB, Resaland GK, Kvalheim OM. Interpretation of Multivariate Association
537 Patterns between Multicollinear Physical Activity Accelerometry Data and Cardiometabolic Health in
538 Children—A Tutorial. *Metabolites.* 2019;9(7):129.

539 33. Kvalheim OM, Arneberg R, Grung B, Rajalahti T. Determination of optimum number of
540 components in partial least squares regression from distributions of the root-mean-squared error
541 obtained by Monte Carlo resampling. *J Chemometrics.* 2018. doi:10.1002/cem.2993.

- 542 34. Rajalahti T, Arneberg R, Berven FS, Myhr KM, Ulvik RJ, Kvalheim OM. Biomarker discovery in mass
543 spectral profiles by means of selectivity ratio plot. *Chemometr Intell Lab Syst.* 2009;95(1):35-48.
544 doi:10.1016/j.chemolab.2008.08.004.
- 545 35. Rajalahti T, Arneberg R, Kroksveen AC, Berle M, Myhr KM, Kvalheim OM. Discriminating Variable
546 Test and Selectivity Ratio Plot: Quantitative Tools for Interpretation and Variable (Biomarker)
547 Selection in Complex Spectral or Chromatographic Profiles. *Anal Chem.* 2009;81(7):2581-90.
548 doi:10.1021/ac802514y.
- 549 36. Cliff DP, Hesketh KD, Vella SA, Hinkley T, Tsiros MD, Ridgers ND et al. Objectively measured
550 sedentary behaviour and health and development in children and adolescents: systematic review and
551 meta-analysis. *Obes Rev.* 2016;17(4):330-44. doi:10.1111/obr.12371.
- 552 37. Rowlands AV, Edwardson CL, Davies MJ, Khunti K, Harrington DM, Yates T. Beyond Cut Points:
553 Accelerometer Metrics that Capture the Physical Activity Profile. *Med Sci Sports Exerc.*
554 2018;50(6):1323-32. doi:10.1249/MSS.0000000000001561.

555 **Figure Legends**

556 **Figure 1. Association patterns between physical activity intensities and a composite**
557 **cardiometabolic health score (ASK dataset) using raw data (min/day), normalized data**
558 **(proportions of wear time), log-transformed data, and compositional data (clr).** The models
559 included 4, 4, 4, and 3 PLS components, respectively. Selectivity ratios are calculated as explained to
560 total variance on the predictive (target projected) component, while retaining the direction of
561 association for each variable

562 **Figure 2. Association patterns between physical activity intensities and locomotor skills (PRESPAS**
563 **dataset) using raw data (min/day), normalized data (proportions of wear time), log-transformed**
564 **data, and compositional data (clr).** All models included 3 PLS components. Selectivity ratios are
565 calculated as explained to total variance on the predictive (target projected) component, while
566 retaining the direction of association for each variable.

567 **Figure 3. Association patterns between physical activity intensities and a composite**
568 **cardiometabolic health score (ASK dataset) using two different descriptions of physical activity**
569 **across the intensity spectrum.** The data (originally described using 23 bins, from 0–99, 100–249,
570 500–999, 1000–1499, ... 9500–9999, to ≥ 10000 cpm, [3]) were condensed to three bins below 4000
571 cpm (left panel) and three bins above 4000 cpm (right panel). Raw data (upper panel), compositional
572 data (clr) (lower panel). Explained variances were 14.34% (3 components) for raw data and 18.37% (3
573 components) for compositional data when the data was condensed below 4000 cpm, and 16.98% (5
574 components) for raw data and 20.76% (5 components) for compositional data when the data was
575 condensed above 4000 cpm. Selectivity ratios are calculated as explained to total variance on the
576 predictive (target projected) component, while retaining the direction of association for each
577 variable.

578

579 **Figure 4. Association patterns between physical activity intensities and locomotor skills (PRESPAS**
580 **dataset) using two different descriptions of physical activity across the intensity spectrum.** The
581 data (originally described using 33 bins, from 0–99, 100–249, 500–999, 1000–1499, ... 14500–14999,
582 to ≥ 15000 cpm, [17]) were condensed to three bins below 4000 cpm (left panel) and three bins
583 above 4000 cpm (right panel). Raw data (upper panel), compositional data (clr) (lower panel).
584 Explained variances were 5.67% (1 components) for raw data and 7.99% (3 components) for
585 compositional data when the data was condensed below 4000 cpm, and 6.35% (2 components) for
586 raw data and 7.96% (2 components) for compositional data when the data was condensed above
587 4000 cpm. Selectivity ratios are calculated as explained to total variance on the predictive (target
588 projected) component, while retaining the direction of association for each variable.

589

590 **Figure 5. Association patterns between physical activity intensities and a composite**
591 **cardiometabolic health score (ASK dataset) using four different log-ratios of physical activity across**
592 **the intensity spectrum.** SEDlr: reference = 0–99 cpm, explained variance = 20.01% (5 components)
593 (upper left panel); LPAIr: reference = 1000–1999 cpm, explained variance = 20.75% (4 components)
594 (upper right panel); MPAIr: reference = 3000–3999 cpm, explained variance = 18.49% (2
595 components) (lower left panel); VPAIr: reference = 7000–7999 cpm (i.e., the intensity strongest
596 associated with cardiometabolic health using raw data), explained variance = 19.90% (5 components)
597 (lower right panel). Selectivity ratios are calculated as explained to total variance on the predictive
598 (target projected) component, while retaining the direction of association for each variable.

599

600 **Figure 6. Association patterns between physical activity intensities and locomotor skills (PRESPAS**
601 **dataset) using four different log-ratios of physical activity across the intensity spectrum.** SEDlr:
602 reference = 0–99 cpm, explained variance = 7.94% (3 components) (upper left panel); LPAIr:
603 reference = 1000–1999 cpm, explained variance = 7.60% (1 components) (upper right panel); MPAIr:

604 reference = 3000–3999 cpm, explained variance = 8.37% (2 components) (lower left panel); VPAI_r:
605 reference = 5000–5999 cpm (i.e., the intensity strongest associated with cardiometabolic health
606 using raw data), explained variance = 8.09% (3 components) (lower right panel). Selectivity ratios are
607 calculated as explained to total variance on the predictive (target projected) component, while
608 retaining the direction of association for each variable.

609

610 **Supplemental Figure 1. Proportions of wear time spent in different PA intensities in the ASK**
611 **dataset (left panel) and PRESPAS dataset (right panel).** Horizontal lines shows the geometric mean
612 composition.

613

614