



1 **Large scale population assessment of physical activity**
2 **using wrist worn accelerometers: the UK Biobank study**

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17

18 **ABSTRACT**

19 **Background:** Physical activity has not been objectively measured in prospective cohorts with
20 sufficiently large numbers to reliably detect associations with multiple health outcomes. Technological
21 advances now make this possible. We describe the feasibility of accelerometer measured physical
22 activity in over 100,000 participants of the UK Biobank study, and its variation by age, sex, day, time of
23 day, and season.

24
25 **Methods:** Participants were approached by email to wear a wrist-worn accelerometer for seven days
26 that was posted to them. Physical activity information was extracted from 100Hz raw triaxial
27 acceleration data after calibration, removal of gravity and sensor noise, and identification of wear /
28 non-wear episodes. We report age- and sex-specific wear-time compliance and accelerometer
29 measured physical activity, overall and by hour-of-day, week-weekend day and season.

30
31 **Results:** 103,712 datasets were received (44.8% response), with a median wear-time of 6.9 days
32 (IQR:6.5–7.0). 96,600 participants (93.3%) provided valid data for physical activity analyses. Vector
33 magnitude, a proxy for overall physical activity, was 7.5% (2.35mg) lower per decade of age (Cohen's
34 $d=0.9$). Women had a higher vector magnitude than men, apart from those aged 45-54yrs. There were
35 major differences in vector magnitude by time of day ($d=0.66$). Vector magnitude differences between
36 week and weekend days ($d=0.12$ for men, $d=0.09$ for women) and between seasons ($d=0.27$ for men,
37 $d=0.15$ for women) were small.

38
39 **Conclusions:** It is feasible to collect and analyse objective physical activity data in large studies. The
40 summary measure of overall physical activity is lower in older participants and age-related differences
41 in activity are most prominent in the afternoon and evening. This work lays the foundation for studies
42 of physical activity and its health consequences. Our summary variables are part of the UK Biobank
43 dataset and can be used by researchers as exposures, confounding factors or outcome variables in
44 future analyses.

45 **INTRODUCTION**

46 Low physical activity is associated with an increased risk of morbidity and mortality [1]. However
47 previous studies are predominantly based on self-reported participation in leisure time activity [2] from
48 which it is difficult to quantify total physical activity across different domains [3]. This uncertainty
49 makes it difficult to convert epidemiological association results into public health recommendations
50 about the minimum level of physical activity required for health and the benefits of engaging in
51 different durations of activity of different intensity. The development of objective methods for assessing
52 physical activity has provided an opportunity to quantify the dose-response relationship of activity with
53 health as a complement to the subjective assessment of self-reported participation in specific
54 activities.

55
56 Accelerometry is the most widely used method for objective assessment of physical activity in
57 population studies [4], and large studies from the UK [5,6], US [7], and Canada [8] indicate age
58 gradients and differences between men and women; time-of-day and day-of-week differences in
59 physical activity. These studies all used accelerometers which were worn around the waist and during
60 awake-time only, a protocol which can result in relatively large amounts of missing data [9]. Therefore,
61 wrist-worn accelerometers are becoming more widely used as an objective measure of physical
62 activity in cohorts in the UK [10], US [9], and Brazil [11]. These devices are water-proof and worn
63 continuously day and night, resulting in higher levels of participant compliance [9,10].

64
65 Cohort studies which include hundreds of thousands of participants followed up over time are required
66 in order to describe the relationship between physical activity and health outcomes that have a
67 number of potential lifestyle, environmental, and genomic causes [12]. Objective assessment of
68 physical activity in such large population-based cohorts has previously not been undertaken because
69 of the challenges of cost and the feasibility of collecting, processing and analysing data on this large
70 scale. In this paper we demonstrate the feasibility of assessing physical activity by wrist-worn
71 accelerometry in the UK Biobank cohort study and report the variation in activity in more than 100,000
72 participants by age and sex.

73

74

75 **METHODS**

76 Study Population

77 UK Biobank is a large prospective study with 500,000 participants aged 40-69 years when recruited in
78 2006-2010 [12]. The study has collected, and continues to collect, extensive phenotypic and genotypic
79 detail about its participants, with ongoing longitudinal follow-up for a wide range of health-related
80 outcomes. Only de-identified data are provided to researchers, who must sign a material transfer
81 agreement, undertaking not to attempt to identify any participant, to keep the data secure, and to use it
82 only for the purposes of the approved research [12]. Between February 2013 and December 2015,
83 participants who had provided a valid email address were sent an email invitation to wear an
84 accelerometer for seven days. The participant email addresses were chosen randomly, with the
85 exception of the North West region which was excluded for much of the project due to participant
86 burden concerns, as this area had been used to trial new projects. From June 2013, participants were
87 sent devices in order of acceptance. This study was covered by the general ethical approval for UK
88 Biobank studies from the NHS National Research Ethics Service on 17th June 2011 (Ref
89 11/NW/0382). None of the authors had direct contact with the study participants.

90

91

92 Accelerometer & Data Collection

93 For objective assessment of physical activity, we used the Axivity AX3 wrist-worn triaxial
94 accelerometer (see **Fig 1**), a commercial version of the Open Movement AX3 open source sensor
95 (<https://github.com/digitalinteraction/openmovement>) designed by Open Lab, Newcastle University.
96 This device demonstrated equivalent signal vector magnitude output on multi-axis shaking tests [13] to
97 the GENEActiv accelerometer used in the Whitehall II [10], Fenland [14] and Pelotas cohorts [11]. The
98 Axivity device facilitates transparent data processing analysis due to its open-source firmware platform
99 and unforced sampling of raw measurement data. We set up the Axivity accelerometers to start at
100 10am two working days after postal dispatch, and capture triaxial acceleration data over a seven day
101 period at 100Hz with a dynamic range of +-8g.

102

103 **Fig 1. UK Biobank triaxial accelerometer and processing steps to extract physical activity**
104 **information.**

105 Axivity AX3 triaxial accelerometer worn on dominant hand as used in UK Biobank (top left). Time
106 series trace of processed accelerometer values after one week of wear (top right). Overview of
107 process to extract proxy physical activity information from raw accelerometer data (bottom).

108

109 Participants were informed in the invitation email and device mail-out letter that the accelerometer
110 should be worn continuously and that they should carry on with their normal activities. Participants
111 were asked to start wearing the accelerometer immediately after receiving it in the post and to wear
112 the monitor on their dominant wrist. They were also informed that the device was configured to
113 automatically turn itself on soon after its arrival and off seven days later. Finally, participants were
114 asked to mail the device back to the co-ordinating centre, in a pre-paid envelope, after the seven day
115 monitoring period.

116

117 Data Processing

118 To ensure different devices provided a similar output under similar conditions we calibrated the
119 acceleration signals to local gravity using the procedure described by van Hees and colleagues [14].
120 Briefly, we identified stationary periods in ten second windows where all three axes had a standard
121 deviation of less than 13.0 mg. These stationary periods were then used to optimise the gain and
122 offset for each axis (9 parameters) to fit a unit gravity sphere using ordinary least squares linear
123 regression. If insufficient data were available to conduct calibration for a given participant (where any
124 of the three sensor axes did not have values outside a ± 300 mg range), we used the calibration
125 coefficients from the previous (or if unavailable, the next) accelerometer record from the same device
126 worn by a different participant. Clipped values, which occur when the sensor's dynamic range of $\pm 8g$
127 is exceeded, were flagged before and after calibration. Recording errors and 'interrupts', which could
128 have occurred for example if participants tried to plug their accelerometer device into a computer,
129 were also logged. Valid data were then resampled to 100 Hz using linear interpolation, except for
130 interrupts lasting longer than 5 seconds which were set to missing. We calculated the sample level
131 Euclidean norm of the acceleration in x/y/z axes, and removed machine noise using a fourth order
132 Butterworth low pass filter with a cutoff frequency of 20Hz. In order to separate out the activity-related
133 component of the acceleration signal, we removed one gravitational unit from the vector magnitude,
134 with remaining negative values truncated to zero [10,11].

135

136 To describe the overall level and distribution of physical activity intensity, we combined the sample
137 level data into five second epochs for summary data analysis, maintaining the average vector
138 magnitude value over the epoch. To represent the distribution of time spent by an individual in
139 different levels of physical activity intensity, we generated an empirical cumulative distribution function
140 from all available five second epochs [11,15]. We removed non-wear time, defined as consecutive
141 stationary episodes lasting for at least 60 minutes where all three axes had a standard deviation of
142 less than 13.0 mg [10,16]. We imputed non-wear data segments using the average of similar time-of-
143 day vector magnitude and intensity distribution data points with one minute granularity on different
144 days of the measurement, as in previous studies [10,16]. This imputation accounts for potential wear
145 time diurnal bias where, for example, if the device was systematically not worn during sleep in an
146 individual, the crude average vector magnitude during wear time would be a biased overestimate of
147 the true average. We then constructed a physical activity outcome variable by averaging all worn and
148 imputed values. Our analysis is freely available and hosted as an open source software project at
149 <https://github.com/activityMonitoring/biobankAccelerometerAnalysis>

150

151 Data Analysis

152 For process evaluation we generated descriptive statistics on the number of participants and devices
153 used. We recorded the number of participants who had insufficient data for calibration. We also noted
154 the percentage of data recording errors caused by interrupts and clipped values, both before and after
155 calibration. Furthermore, we described the number of participants who provided different amounts of
156 wear time. We then excluded individuals with less than three days (72 hours) of wear data or who did
157 not have wear data in each one-hour period of the 24-hour cycle. We defined this criteria after finding
158 72 hours of wear were needed to be within 10% of a complete seven day measure (using intraclass
159 correlation coefficients) in missing data simulations on 29,765 participants who had perfect wear time
160 compliance (see S1 – minimum wear time criterion).

161

162 Descriptive statistics were used to report device wear time compliance in hours and accelerometer
163 measured physical activity in milli-gravity units (mg). Age groups were categorised into decade bands
164 from ages 45-79 years. Age and seasonal (with Spring starting on 1st March) differences in device

165 wear-time were examined using the Kruskal-Wallis test, while sex differences were examined using
166 the Wilcoxon-Mann Whitney test. Differences in wear-time distribution were examined using the
167 Friedman test for time-of-day (six hour quadrants, e.g. 00:00-05:59, 06:00-11:59, etc.) and Wilcoxon
168 signed ranks test for days (weekdays versus weekend days), within individuals for men and women
169 separately. Mean acceleration vector magnitude differences by age group were investigated using
170 one-way repeated measures ANOVA for time-of-day (six hour quadrants) and days (weekdays versus
171 weekend days), within individuals for men and women separately. Seasonal differences in mean
172 acceleration vector magnitude were investigated using two-way ANOVA between age groups, for men
173 and women separately. We used R to perform all statistical analysis [17]. Given the size of this
174 dataset, almost all of our findings show robust statistical significance ($p < 0.001$). We therefore do not
175 report such small p-values.

176

177

178 **RESULTS**

179 A total of 236,519 UK Biobank participants were approached, of whom 106,053 agreed to wear a
180 physical activity monitor (44.8%). The median time between each participant being invited to take part
181 and being sent a device was 113 days (IQR: 73 – 137 days). Fig 2 shows that 103,712 datasets were
182 received for data analysis. 123 participants were excluded as they were aged less than 45 years.
183 Eleven participants were excluded from further analysis; eight because the calibration by the
184 preceding or subsequent measurement was not possible due to insufficient data; and three
185 participants due to unreliable device data. A total of 4043 devices were used on a median number of
186 27 occasions (IQR: 8 – 39). The median time between each device being posted was 17.0 days (IQR:
187 15.8 – 19.8) with a median of 832 devices (IQR: 629 – 994) posted each week.

188

189 **Fig 2. Participant flow chart; the UK Biobank study 2013-2015 (n= 103,712).**

190

191 Calibration of the data to local gravity greatly reduced the error in the assessment of acceleration with
192 the root mean square error of stationary points falling from an average of 81.8 mg (95% CI: 81.6 –
193 82.1) to an average of 2.6 mg (95% CI: 2.6 – 2.6). However, 2.9% (n=3049) of participants had
194 insufficient stationary data to inform the calibration. These individual records were calibrated using
195 stationary episodes from the previous (n=2887) or next (n=154) use of the same device by different

196 participants. The influence of clips (readings beyond the sensor's dynamic range of +8g) before
 197 (median: 160, IQR: 62 – 393) and after (median: 169, IQR: 67 – 410) calibration, interrupts (median: 0,
 198 IQR: 0 – 0), and errors such as clips or missing readings (median: 200, IQR: 66 – 355) was negligible,
 199 with respect to the median of 58.6 million data readings (IQR: 56.0 – 60.1 million).

200
 201 Fig 3 illustrates that 80.6% of participants wore the device for at least 150 hours out of a scheduled
 202 168 hours. Men wore the device for a median of 166.3 hours (IQR: 157.7 – 168.0) and were slightly
 203 more compliant than women who wore the device for a median of 165.6 hours (IQR: 156.7 – 167.0).
 204 Table 1 shows that older age groups had marginally higher levels of compliance than younger age
 205 groups. Analysis of wear time compliance by age on a linear scale shows that on average there was a
 206 difference of 2 hours 18 minutes (1.6%) for each decade. In addition, Table 1 indicates minimal
 207 differences in the wear time compliance by time-of-day and week-weekend day. No wear-time
 208 differences were found by season. We removed 6978 (6.7%) participants who had insufficient wear
 209 data for our remaining analyses on accelerometer measured physical activity.

210

211 **Table 1. Wear-time compliance and acceleration vector magnitude by age, day, time of day, and**
 212 **season, stratified by sex: The UK Biobank study 2013-2015 (n=103,578).**

Age (yrs) ^A	Wear time [median (IQR) hours]		Acceleration vector magnitude [mean +- stdev mg]	
	Women	Men	Women	Men
45-54	164.9 (152.4 – 167.0) (n=12,586)	165.4 (149.5 – 168.0) (n=8655)	31.2 +- 8.7 (n=11,572)	31.1 +- 9.7 (n=7838)
55-64	165.4 (156.0 – 167.0) (n=21,322)	165.8 (156.5 – 168.0) (n=14,410)	29.1 +- 8.0 (n=19,890)	28.8 +- 8.8 (n=13,362)
65-74	165.6 (159.1 – 168.0) (n=22,821)	166.8 (160.8 – 168.0) (n=20,595)	26.6 +- 7.1 (n=21,489)	25.6 +- 7.7 (n=19,385)
75-79	165.6 (158.9 – 167.0) (n=1494)	166.8 (162.6 – 168.0) (n=1695)	23.9 +- 6.5 (n=1436)	22.9 +- 6.8 (n=1628)
p value	p<0.001	p<0.001	p<0.001	p<0.001
Time of day^B				
0 - 5.59 _{am}	40.9 (36.0 – 42.0) (n=58,223)	42.0 (36.6 – 42.0) (n=45,355)	4.4 +- 3.1 (n=54,387)	4.9 +- 4.4 (n=42,213)
6 - 11.59 _{am}	41.0 (38.9 – 42.0) (n=58,223)	42.0 (39.0 – 42.0) (n=45,355)	38.6 +- 14.9 (n=54,387)	37.4 +- 16.4 (n=42,213)
12 - 5.59 _{pm}	42.0 (40.3 – 42.0) (n=58,223)	42.0 (40.3 – 42.0) (n=45,355)	44.3 +- 13.8 (n=54,387)	42.9 +- 16.0 (n=42,213)
6- 11.59 _{pm}	42.0 (39.5 – 42.0) (n=58,223)	42.0 (40.2 – 42.0) (n=45,355)	26.4 +- 10.4 (n=54,387)	24.9 +- 11.5 (n=42,213)
p value	p<0.001	p<0.001	p<0.001	p<0.001
Day^C				
Weekday	23.7 (22.5 – 24.0)	23.8 (22.6 – 24.0)	28.5 +- 8.2	27.5 +- 9.0

	(n=58,223)	(n=45,355)	(n=54,387)	(n=42,213)
Weekend	24.0 (22.9 – 24.0)	24.0 (23.3 – 24.0)	28.0 +- 9.4	27.1 +- 10.8
	(n=58,223)	(n=45,355)	(n=54,387)	(n=42,213)
p value	p<0.001	p<0.001	p<0.001	p<0.001
Season^D				
Spring	165.6 (156.2 - 167.5)	166.1 (157.4 - 168.0)	28.8 +- 8.0	28.1 +- 9.1
	(n=13,365)	(n=10,224)	(n=12,480)	(n=9,469)
Summer	165.4 (156.2 - 168.0)	166.3 (157.4 - 168.0)	28.8 +- 8.1	28.2 +- 8.7
	(n=15,450)	(n=11,943)	(n=14,353)	(n=11,016)
Autumn	165.6 (157.2 - 167.0)	166.3 (158.2 - 168.0)	28.3 +- 8.0	27.3 +- 8.7
	(n=17,213)	(n=13,506)	(n=16,157)	(n=12,633)
Winter	165.6 (156.7 - 168.0)	166.3 (157.9 - 168.0)	27.7 +- 7.8	26.3 +- 8.4
	(n=12,195)	(n=9,682)	(n=11,397)	(n=9,095)
p value	p=0.289	p=0.104	p<0.001	p<0.001

213

214 ^A Age: Kruskal-Wallis test used to compare wear-time distributions, and one-way analysis of variance test used to compare
215 acceleration vector magnitude means. Sum wear time hours for week displayed (max = 168.0).

216 ^B Time of day: Friedman test used to compare wear-time distributions within individuals, and repeated one-way analysis of
217 variance test used to compare acceleration vector magnitude means within individuals and between age groups. Sum wear time
218 hours for time quadrant over a week displayed (max = 168.0).

219 ^C Day: Wilcoxon test used to compare wear-time distributions within individuals, and repeated one-way analysis of variance test
220 used to compare acceleration vector magnitude means within individuals and between age groups. Average wear time hours for
221 day displayed (max = 24.0).

222 ^D Season (Spring starting on 1st March): Kruskal-Wallis test used to compare wear-time distributions, and two-way analysis of
223 variance test used to compare acceleration vector magnitude means between age groups. Sum wear time hours for week
224 displayed (max = 168.0).

225

226 **Fig 3. Cumulative distribution function of accelerometer wear time compliance; the UK Biobank**
227 **study 2013-2015 (n= 103,578).**

228

229 Table 1 describes the variation in mean vector magnitude, the summary measure of accelerometer
230 measured physical activity, by age and sex in the sub-group of 96,600 participants who had good
231 wear time compliance. Vector magnitude was higher in women than men, apart from those aged 45-
232 54 years (p = 0.98). The mean effect size for these sex differences was small (0.09), ranging from
233 0.01 for 45-54 years to 0.15 for 75-79 years. There was strong evidence of accelerometer measured
234 physical activity differing by age group in both men and women. The mean physical activity in the age
235 group 45-54 years was 31.17 mg (SD 9.10) and was, on average 7.5% or 2.35 mg lower per decade.
236 The mean effect size for these age differences was large, at 0.89 for women and 0.9 for men. Fig 4
237 shows the distribution of the data within age and sex strata, highlighting that although there appears to

238 be an overall decline in average physical activity with increasing age, there is considerable overlap in
239 the distributions with many older participants being more active than those in the youngest age
240 category.

241

242 **Fig 4. Acceleration vector magnitude by sex and age; the UK Biobank study 2013-2015 (n=**
243 **96,600).**

244

245 Fig 5 shows the mean physical activity level by hour of day averaged across the whole measurement
246 period by age and sex. It shows that the effect size for physical activity differences between age
247 groups are most apparent in the afternoon (0.74 for women and 0.69 for men) and evening (1.06 for
248 women, 1.12 for men) with smaller differences by age group in the morning (0.56 for women, 0.46 for
249 men). Weekdays and weekend days differed, with vector magnitude higher at weekdays except for
250 those aged 45-54 years. However, the mean effect size for these day differences was small (0.10),
251 ranging from 0.04 to 0.15 across female age groups and 0.11 to 0.18 for male age groups (see Fig 6).
252 Seasonality also differed, with vector magnitude lower during winter months except for women aged
253 75-79. However, the mean effect size for these season differences was small (0.21), ranging from
254 0.09 to 0.18 across age groups in women and 0.17 to 0.41 across age groups in men (see Fig 6).

255

256 **Fig 5. Variation in mean acceleration across the day by age and sex: the UK Biobank study**
257 **2013-2015 (n= 96,600).**

258 Shading bounds represent two standard errors.

259

260 **Fig 6. Acceleration vector magnitude by day of the week (top), season (bottom), age, and sex:**
261 **the UK Biobank study 2013-2015 (n= 96,600).**

262

263 To illustrate time spent at different physical activity intensities, Fig 7 plots the empirical cumulative
264 distribution function of the five second sample values for each subgroup. The bottom part of this figure
265 shows sex differences in the distribution of physical activity intensity, for each age group. For example,
266 men spend more time at or below 25 mg than women (122.6 versus 119.3 hours), but also slightly
267 more time above 225 mg than women (2.18 versus 2.09 hours).

268

269 **Fig 7. Cumulative time spent in various acceleration categories by sex and age (top), and sex**
270 **differences by age and intensity level (bottom); the UK Biobank study 2013-2015 (n=96,600)**

271

272

273 **DISCUSSION**

274 Developments in the technology supporting objective assessment of physical activity have now made
275 it possible to consider assessing this behaviour objectively in large scale population-based cohort
276 studies as an adjunct to more traditional assessment of self-reported participation in activities within
277 different domains of life. However, even with those technological developments, it has previously been
278 unknown whether it would be possible for this approach to be acceptable to participants and whether it
279 would prove to be feasible to collect, analyse and interpret data from over one hundred thousand
280 participants. This report from the UK Biobank study shows that 45% of participants who were invited to
281 wear a monitor accepted the invitation. It also shows that measuring activity with a wrist worn device is
282 highly acceptable to participants as manifest by the very high proportion of people in whom the data
283 were of high quality and completeness. By necessity in the UK Biobank Study participants were invited
284 to wear the monitor some time after recruitment to the baseline visit. As with all add-on measurements
285 that are conducted on a different occasion, there will be participants who don't accept the invitation to
286 participate. Other studies in which wrist worn accelerometers are part of the protocol for a baseline
287 visit, rather than a separate add-on, will be likely to achieve higher participation levels.

288

289 We have shown that mean vector magnitude in this population was greater in women than men, apart
290 from those aged 45-54 years. Whether these results indicate true differences in physical activity
291 between sexes or are a function of a between-sex difference in the relationship between wrist
292 acceleration and true activity remains to be investigated. In addition, we observed a marked overall
293 difference in the summary measure of physical activity by age, with older participants having levels of
294 activity that are, on average, 7.5% lower for each 10 year age difference. These differences by age
295 group are similar to other population-based studies [5,7] that have used hip worn accelerometers. Our
296 findings also suggest that men spend more time than women in what might be considered low or
297 sedentary levels of physical activity, while women spend more time in moderate levels of activity.

298 Older participants are much less active than younger participants during afternoons/evenings than in
299 the morning, which mirrors previous findings in older UK adults using hip worn accelerometers [6].
300 There were small differences between weekday and weekend day physical activity, and also small
301 seasonal differences in activity. We have not generalised the overall descriptive findings to the UK
302 population since the UK Biobank was established as an aetiological study rather than one aimed at
303 population surveillance [5,7].

304
305 We have extracted objective physical activity information from 103,578 participants aged 45-79, who
306 were asked to wear accelerometers for seven days on their dominant wrist. The strengths of this study
307 include its use of objective measures of physical activity, excellent participant compliance,
308 unprecedented scale, and use of reproducible methods. For example, >93% of participants provided
309 more than 72 hours of wear time with no missing data bias by time of day. The overall levels of
310 participant compliance in the UK Biobank mirror findings in other studies that have used wrist-worn
311 accelerometers in thousands of participants [9–11]. Wrist-worn accelerometers are not only highly
312 acceptable to participants, but are also valid. Laboratory-based studies have demonstrated that the
313 signal from wrist-worn devices correlates with physical activity energy expenditure as well as
314 traditional waist-worn devices ($R=0.83$ vs. $R=0.87$) [18]. However, more robust validation studies of
315 physical activity information from raw wrist-worn accelerometer data are needed to enhance the
316 interpretation of this signal.

317
318 Even though we used relatively simple summary measures in these analyses, their derivation still
319 involved several critical data processing decisions, the alteration of which would have large effects on
320 the derived physical activity variables [14]. For example, there is uncertainty on how to address
321 negative values during the gravity removal process. Furthermore, it is not possible to perfectly
322 separate static and dynamic acceleration (for example gravity and physical activity) from the
323 measurement of triaxial acceleration alone. Therefore, we produced summary statistics of vector
324 magnitude which do not attempt this separation. In addition, absolute and truncated Euclidian norm
325 minus one and high-pass filtered vector magnitude (all of which attempt to separate out activity) were
326 generated too. We found that while the magnitude of these variables changes, their correlation was
327 very strong (>0.95), which provides confidence in our chosen metric for association studies.

328 Uncertainty also exists on the best method to identify non-wear episodes, and the size of epoch on
329 which to base distributions of physical activity intensity. While the derived factors are only the most
330 basic variables that can be extracted from the raw 100Hz triaxial acceleration data, future projects will
331 be able to build on this foundation to derive additional parameters describing other aspects of physical
332 activity, sedentary behaviour and sleep.

333

334 In conclusion, the collection and processing of this large accelerometer dataset in a prospective cohort
335 study lays the foundation for studies of physical activity and its health consequences. The summary
336 variables that we have constructed are now part of the UK Biobank dataset and can be used by
337 researchers as exposures, confounding factors or outcome variables in future analyses.

338

339

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403 **SUPPORTING INFORMATION**

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405 **S1 – Minimum wear time criterion**













