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1 **Jumping in the night: an investigation of leaping activity of western tarsier**
2 **(*Cephalopachus bancanus borneanus*) using accelerometers**

3

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23

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25 **Abstract**

26 Accelerometers enable scientists to quantify activity of free-living animals whose direct
27 observation is difficult or demanding due to their elusive nature or nocturnal habits.
28 However, the deployment of accelerometers to small-bodied animals and, in particular,
29 to primates has been little explored. Here we show the first application of
30 accelerometers on the Western tarsier (*Cephalopachus bancanus borneanus*), a
31 nocturnal small-bodied primate endemic to the forests of Borneo. The fieldwork was
32 carried out in the Lower Kinabatangan Wildlife Sanctuary, Sabah, Malaysian Borneo.
33 We provide guidelines for the deployment of accelerometers on tarsiers that might also
34 be applied to other primate species. Our data collected on two females show levels of
35 leaping activity comparable to those previously described using direct observation of
36 wild or captive individuals. The two females showed different patterns of leaping
37 activity, which calls for work to explore individual differences further. Our work
38 demonstrates that accelerometers can be deployed on small primates to collect body
39 motion data that otherwise would be demanding to collect using classic field
40 observations. Future work will be focused on using accelerometer data to discriminate
41 in more detail the different behaviours tarsiers can do and to address the causes and
42 consequences of individual variation in activity.

43

44 **Key Words**

45 Body acceleration · Movement ecology · Primates · Tarsier

46

47 **Running head:** Accelerometers and tarsiers

48

49

50 **Introduction**

51 Activity patterns can provide important information about the ways animals interact
52 with their environment. Historically, however, this type of data has been not only time-
53 consuming but also logistically challenging to collect, particularly for elusive or
54 nocturnal species. In recent years, monitoring activity with accelerometer loggers is
55 increasingly being used as accelerometers can remotely record the body motions of
56 free-living animals [e.g., Sullivan et al., 2006; Byrnes et al., 2011; Nathan et al., 2012;
57 Wunderlich et al., 2014; Hammond et al., 2016]. Accelerometer loggers are small
58 devices that can be attached to an animal's body (e.g., on a collar or backpack) and take
59 recordings of either bi- or tri-axial body acceleration. Gravitational acceleration
60 produces a signal referred to as static acceleration, which depends on the posture of the
61 tagged individual (when the accelerometer is fixed to the subject). The signal derived
62 from the subject's motion is referred to as dynamic acceleration. Data on dynamic
63 acceleration can be used to calculate the vector sum of dynamic body acceleration
64 (VeDBA) or the overall dynamic body acceleration (ODBA), which indicate body
65 motion when g-values are higher than 0. Both VeDBA and ODBA values may also
66 provide information about energy expenditure [e.g., Qasem et al., 2012; Jeanniard-de-
67 Dot et al., 2017]. Accelerometer data can be used to derive daily activity budgets [e.g.,
68 when an animal moves; Yoda et al., 1999; Lagarde et al., 2008; Gr unew lder et al.,
69 2012; Williams et al., 2014] and, if accelerometer and viewer-observed behavioural data
70 are collected simultaneously, accelerometer data can also be used to characterise
71 behaviours (e.g., walking, running, leaping [Sakamoto et al., 2009; Nathan et al., 2012;
72 Graf et al., 2015] and to estimate energy expenditure associated with different
73 behaviours [Wilson et al., 2006; Qasem et al., 2012; Jeanniard-du-Dot et al., 2016].

74 Until now, accelerometers have primarily been deployed on large-bodied species [Fig. 1
75 in Hammond et al., 2016], but with the development of micro-accelerometers, the
76 deployment of accelerometers on small-bodied animals is increasing rapidly [Hammond
77 et al., 2016].

78 In recent times, there has been growing interest of primatologists in
79 accelerometers, but the number of studies carried out on free-living animals is still
80 limited [Sellers and Crompton, 1994, 2004; Mann et al., 2005; Sullivan et al., 2006;
81 Papailiou et al., 2008; Ravignani et al., 2013; Wunderlich et al., 2014]. Prior work on
82 primates found that accelerometers can enable investigators to collect valuable
83 information about the behaviour and ecology of a given species. For example, Sullivan
84 et al. [2006] found that the level of physical activity quantified using accelerometers is a
85 particularly important factor contributing to weight change in adulthood and that there
86 are large, but stable, differences in physical activity among individuals in female rhesus
87 monkeys (*Macaca mulatta*). Wunderlich et al. [2014] showed that data collected with
88 accelerometers can enable the identification of specific movement patterns of
89 Verreaux's sifakas (*Propithecus verreauxi*) in the absence of direct observation.
90 Moreover, accelerometers might offer primatologists and conservation biologists a tool
91 to investigate how environmental changes or even captive care techniques can affect
92 individual activity.

93 Western tarsiers are small primates endemic to the tropical forests of Borneo and
94 are currently listed as Vulnerable by IUCN. Their nocturnal behavioural patterns and
95 small size makes direct visual observation of them demanding. Previous work on both
96 captive and wild tarsiers showed that they have a bimodal nocturnal activity (mostly due
97 to leaping), with peaks of activity occurring shortly after sunset and slightly before

98 sunrise (Niemitz, 1984; Crompton and Andau, 1986, 1987). In this study, we deployed
99 accelerometers on the Western tarsier (*Cephalopachus bancanus borneanus*) for the
100 first time and used this earlier work on tarsiers as a reference to compare our
101 accelerometer data. Here we provide guidelines for the deployment of accelerometers
102 on tarsiers and assess whether the data gathered by accelerometers can deliver valuable
103 information on leaping activity, which is the main locomotor mode of tarsiers (more
104 than 60% in Crompton and Andau, 1986), that would otherwise be demanding to collect
105 using traditional field observations (Crompton and Andau, 1986, 1987).

106

107 **Materials and Methods**

108 The fieldwork was carried out from the 1st to the 21st of April 2016 in the Lower
109 Kinabatangan Wildlife Sanctuary, East Sabah, Malaysian Borneo. The entire sanctuary
110 spans 26,100 ha along both sides of the Kinabatangan River. Data were collected within
111 the areas surrounding the Danau Girang Field Centre (Lot 6, N5° 24' 49.4" E118° 02'
112 14.9"), a collaborative research and training facility managed by the Sabah Wildlife
113 Department and Cardiff University.

114 Body acceleration was measured using small (size of the whole unit: 0.8×10×19
115 mm; weight of the whole unit plus shrinkable plastic tube: 1.2 g, ≤ 1% of a tarsier body
116 mass), ultra-low power, 3-axis acceleration data loggers (AXY-3, Technosmart Europe
117 srl, Roma, Italy). The AXY Manager software (Technosmart Europe srl, Roma, Italy)
118 was used to configure the accelerometers and download the data. Each device was
119 connected to a laptop using a USB cable and configured to record continuously at a
120 frequency of 10 Hz (i.e., 10 readings taken per second) with a 10-bit sampling
121 resolution and a g-range of ± 4. We chose a frequency of 10 Hz because at this

122 sampling rate our accelerometers can record continuously the body motion for over one
123 month. Prior work on other species used a sampling rate of 100 Hz [Byrnes et al., 2011;
124 Wunderlich et al., 2014], but this required the application of heavier accelerometers for
125 shorter periods than ours. However, a sampling rate of 10 Hz is low and might not
126 capture all leaps, hence we configured the accelerometers in order to have a sampling
127 resolution of 10 bits. Accelerometers were deployed on two adult females (herein F1
128 and F2, both with a body mass of 135 g) that had previously been radio-collared. At the
129 time of accelerometer deployment, each female had a three-month old infant. The
130 animals were located in the forest during the day using radio telemetry; once spotted,
131 they were captured by hand and put in a cotton bag to keep them calm. Once a tarsier
132 was caught, the radio collar (Biotrack Ltd PIP3 Tag with coated brass collar; 3.55g) was
133 removed to attach the accelerometer with electrical tape, then was covered with a
134 transparent shrinkable plastic tube to provide abrasion resistance and environmental
135 protection (fig. 1). Using a lighter, the plastic tube was slightly melted to make it adhere
136 to the accelerometer. When doing so, the lighter had to be moved continuously along
137 the plastic tube to avoid a concentration of heat in a given spot because, while
138 accelerometers are resistant to heat, cables (that connect the battery to the
139 accelerometer) and the battery could be damaged. The accelerometer was attached on
140 the radio-collar in such a way that made it stay on the dorsal side of the head during the
141 study period. We opted to attach the accelerometer to the radio-collar because
142 accelerometers had to be recollected in order to download the data onto a computer, and
143 radio-collars allowed us to easily locate the tarsiers again. Moreover, the radio-collar
144 provided a support where to attach the accelerometer without the need to use other
145 techniques, which helped to keep the overall extra-weight low. Before releasing the

146 animal, the accelerometer was turned on by passing a magnet over the magnetic switch
147 near the connector area of the device. Successful activation of the accelerometers was
148 signalled by the blinking blue light indicator, which then switched off after 29 flashes.
149 We recorded the time at which the accelerometer was turned on, which is needed in
150 order to link accelerometer data to the time of day they were recorded. When tarsiers
151 were re-caught, the accelerometers were turned off by using the same magnet as that
152 used to turn them on, and then removed from the collar. When back to the field station,
153 accelerometers were connected to a laptop using a USB cable and the AXY Manager
154 software was used to download the data recorded by the accelerometers.

155 The Framework4 software (<http://framework4.co.uk/index.php>) was used to
156 visualise and analyse the accelerometer data that had been downloaded with the AXY
157 Manager software. The raw values of acceleration for each X, Y and Z axis were the
158 result of the combination of static acceleration (due to gravity) and dynamic
159 acceleration (due to movement), and were derived using the software. Raw
160 accelerometer data were converted to dynamic body acceleration by first using a
161 moving average to smooth each channel in order to derive the static acceleration and
162 then subtracting this static acceleration from the raw data. The gravitational component
163 (which sums to one) is therefore already subtracted before the calculation of VeDBA.
164 Using the same software, the vector sum of dynamic body acceleration in three
165 dimensions [$\text{VeDBA} = \sqrt{A_x^2 + A_y^2 + A_z^2}$] was calculated and used as an index of body
166 motion [Qasem et al., 2012]. Although accelerometers were well attached on the radio-
167 collar, VeDBA instead of ODBA was used because, conversely to the ODBA, the
168 VeDBA values are less affected by any inconsistent orientation of the devices among
169 individuals.

170

171 **Results**

172 **Performance of accelerometers**

173 Data on free-ranging animals were collected for 78 and 311 hours for F1 and F2,
174 respectively. Shortly after deployment, the accelerometer of F1 unexpectedly stopped
175 recording, probably due to battery failure. A possible reason may be due to insufficient
176 waterproofing during deployment, resulting in damage to the battery during a heavy
177 rainstorm three days after deployment. The battery of F2 was substantially more than
178 half full when we re-caught the tarsier. Thus, with our configuration, body acceleration
179 data may be collected for around 35-40 days. Fig. 2A shows the overall acceleration
180 profile of F1, while fig. 2B shows how the number of peaks of the acceleration profile
181 increases during the transition from dormancy to arousal for F1.

182

183 **Identification of leaps**

184 Fig. 3 shows the acceleration signals of F2 during the recapture session; the tarsier
185 jumped away repeatedly before successfully re-capturing her. Our recording with a
186 video camera of F2 enabled us to ascertain that only leaps generated VeDBA values at
187 least ≥ 1 g (fig. 3). Using the Framework4 software, we could link accelerometer data to
188 the time of the day at which a given accelerometer value was recorded. Thus we could
189 link the accelerometer values to the video recording by matching the video to the time at
190 which a leap was made. During our video recording, we also observed tarsiers to climb
191 and to turn their head in different directions. We also observed the two tarsiers climbing
192 and turning their head after being released after accelerometer deployment. In all these
193 observations, these behaviours did not generate VeDBA values higher than 1. However,

194 VeDBA values below 1 g might also indicate small leaps. To be conservative, a cut-off
195 VeDBA value of 1 was deemed appropriate to estimate the number of leaps made by
196 each tarsier and to describe leaping activity in this study. However, given the limited
197 duration of our direct observations (less than one hour), we recommend that future
198 protocols would include longer recordings of the animals to make sure that all
199 behaviours are observed and the accelerometer response to them tested.

200

201 **Comparison of leaping activity**

202 To make data of F1 and F2 comparable, we only took the first 78 hours of recording for
203 F2 into account. Our data show that F1 shows less leaping activity (i.e., number of leaps
204 made) than F2 during the night (fig. 4). The main period of leaping activity for the two
205 tarsiers began between 17:00 and 18:00 and ended between 05:00 and 06:00. For both
206 tarsiers, there was a peak of leaping activity between 18:00 and 19:00 (black arrow in
207 fig. 4), but for F2 there were additional peaks of leaping activity between 21:00 and
208 22:00 and between 04:00 and 05:00 (black arrow in fig. 4). For F1, we found low
209 nocturnal leaping activity between 20:00 and 06:00, whereas the period of low
210 nocturnal leaping activity was only between 00:00 and 02:00 for F2 (fig. 4). As
211 expected, both tarsiers were most active in terms of leaps made during the night,
212 however, there were also values of VeDBA higher than 1 (mainly above 3) during the
213 day, mainly between 10:00 and 11:00 and between 12:00 and 13:00 for F1 (grey arrow
214 in fig. 4); between 12:00 and 14:00 for F2 (grey arrow in fig. 4). The leaping activity of
215 F2 recorded over the selected period of 78 hours was very similar to that recorded over
216 the entire period, i.e. 311 hours (fig. 5). The number of leaps made by F2 per night is
217 similar to that estimated by previous work on tarsiers, while that of F1 was much

218 smaller (fig. 6).

219

220 **Discussion**

221 Our results show that accelerometers can be used successfully to collect data on body
222 motion that can be used to quantify the leaping activity of tarsiers. It is, however, very
223 important to protect the devices to avoid any damage caused by the animal or
224 environment. This might be particularly relevant if accelerometers are deployed on
225 primates that live in groups, where engaging in social grooming can lead to removal or
226 destruction of the unit. Building upon previous studies on primates in captivity [Sellers
227 and Crompton, 1994, 2004; Mann et al., 2005; Sullivan et al., 2006; Papailiou et al.,
228 2008; Ravignani et al., 2013; Wunderlich et al., 2014], our study shows that
229 accelerometers can also be deployed on free-living primates as small as a tarsier for
230 several weeks.

231 Our data on leaping activity of tarsiers are generally in agreement with previous
232 work on the activity of tarsiers. A previous study using a continuous all-night following
233 of free-living tarsiers concluded that the Western tarsier is entirely nocturnal [Crompton
234 and Andau, 1986, 1987]. Tarsiers began to move from their sleeping sites between
235 18:05 and 19:10 but most often between 18:30 and 18:45, and would stop traveling
236 between 05:55 and 06:15 [Crompton and Andau, 1987]. Previous work also showed that
237 the nocturnal leaping activity of tarsiers appeared to be bimodal, with peaks of activity
238 shortly after sunset and slightly before sunrise. Niemitz [1984] found a peak in leaping
239 activity in captive tarsiers around 19:00 and then a second peak between 05:00 and
240 06:00. Crompton and Andau [1987] found similar results in wild tarsiers. A study on
241 captive individuals found that tarsiers were almost completely sedentary during the

242 photoperiod (inactive period), with changes in location recorded on only 5% of 1,576
243 position checks during the day on 408 randomly selected days [Roberts and Kohn,
244 1993]. Tarsiers generally awoke within 15 minutes of the onset of the night, and
245 virtually all activity occurred during the night [Roberts and Kohn, 1993]. Our estimates
246 of leaping activity may be considered conservative because we do not know if VeDBA
247 values lower than 1 indicate small leaps (a g-value of 2 g is equivalent to a leap of
248 approximately 1 m in the red ruffed lemur (*Varecia rubra*); Sellers and Crompton,
249 2004) and our low sampling rate (10 Hz) might have not captured all the leaps made.
250 However, the estimated number of leaps made by F2 per night is similar to that
251 estimated by previous studies on both captive and wild tarsiers using direct observations
252 of the animals (fig. 5). The number of leaps made by F1 was lower, reasons of which
253 need further exploration. Our data show that tarsiers may occasionally leap during the
254 daytime, the reasons of which need to be further explored. Previous work on primates
255 found that factors, such as human disturbance, personality or availability of food [e.g.,
256 Krebs and Davies, 1993; Peres, 1993; Passamani, 1998; Uher et al., 2008], may
257 influence individual activity budgets. Studies on other species, such as the Malayan
258 colugos (*Galeopterus variegatus*), also showed that sexes may differ in activity budgets
259 and performance of different locomotor behaviours [Byrnes et al., 2011]. Furthermore,
260 we cannot exclude the possibility that the two tarsiers responded differently to the
261 deployment of accelerometers. More research is necessary to address this issue (e.g.,
262 testing longer lasting deployments, assessing metrics of physiological stress).

263 In conclusion, accelerometers enabled data collection on leaping activity without
264 the need of doing continuous all-night follows. However, this study did not enable us to
265 determine specific behaviours, such as if leaping indicated foraging or fleeing from a

266 threat. To do so, controlled observations of tarsiers in the wild or in captivity are
267 required to link acceleration data to specific behaviours. Future work will be needed to
268 assess the extent to which accelerometer data can be used to determine specific
269 behaviours of tarsiers. In doing so, it will be important to test whether recording at
270 frequencies higher than that used in this study (i.e., 10 Hz) would improve
271 differentiation between behaviours and estimate of the number of leaps. Understanding
272 more about the behaviour of tarsiers can have important implications for their
273 conservation. For example, data on daily activity may give information on the impact of
274 human disturbance or quality of the environment (e.g., tarsiers might move more for
275 foraging in forest patches near plantations) has on tarsiers. Moreover, validation of
276 VeDBA [or of ODBA; Jeanniard-de-Dot et al., 2017] as a metric of energy expenditure
277 might provide a tool to investigate the metabolic costs incurred due to the rapid land-use
278 changes that are occurring in the tropics.

279

280 **Conflict of interest statement**

281 The authors certify that they have no any conflicts of interest.

282

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295

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377

378 **Figure captions**

379 Figure 1. Example of how the tri-axial accelerometer was attached on the radio-collar
380 deployed on a Western tarsier (*Cephalopachus bancanus borneanus*).

381

382 Figure 2. The accelerometer profile of Western tarsier (*Cephalopachus bancanus*
383 *borneanus*) F1 (Panel A) and the accelerometer profile of F1 during the transition from
384 dormancy to arousal (Panel B).

385

386 Figure 3. Accelerometer profile of the Western tarsier (*Cephalopachus bancanus*
387 *borneanus*) F2 that was video-recorded while leaping away during an attempted re-
388 capture. The VeDBA peaks shown in the graph refer only to when the tarsier was seen
389 leaping. Acc. X = acceleration along the X axis; Acc. Y = acceleration along the Y axis;
390 Acc. Z = acceleration along the Z axis; VeDBA = vector sum of dynamic body
391 acceleration in the three dimensions.

392

393 Figure 4. Leaping activity of the two female Western tarsiers (*Cephalopachus bancanus*
394 *borneanus*) over a period of 78 hours. Note that only VeDBA-values ≥ 1 , which
395 indicate leaping behaviour, were used for this description; thus data are not
396 representative of the whole activity budgets. Note also that we are using counts, which
397 refer to the peaks of the VeDBA profile. Our accelerometers recorded ten times per
398 second, meaning that a single leap included several g-values, which increased as soon as
399 the animal jumped and then decreased as soon as the animal started landing. Values are
400 shown as mean and standard deviation per hour. Grey dots refer to F1, while black dots
401 refer to F2. Note that in order to avoid overlap of the boxplots of the two tarsiers,

402 boxplots of F2 were slightly moved to the right of those of F1. Grey arrows indicate
403 leaping activity during the daylight, while black arrows indicate peaks of leaping
404 activity during the night.

405

406 Figure 5. Leaping activity of one female Wester tarsier (*Cephalopachus bancanus*
407 *borneanus*) recorded over a period of 311 hours.

408

409 Figure 6. Comparisons of estimated average number of leaps made per night by tarsiers
410 recorded in our and previous studies.

411

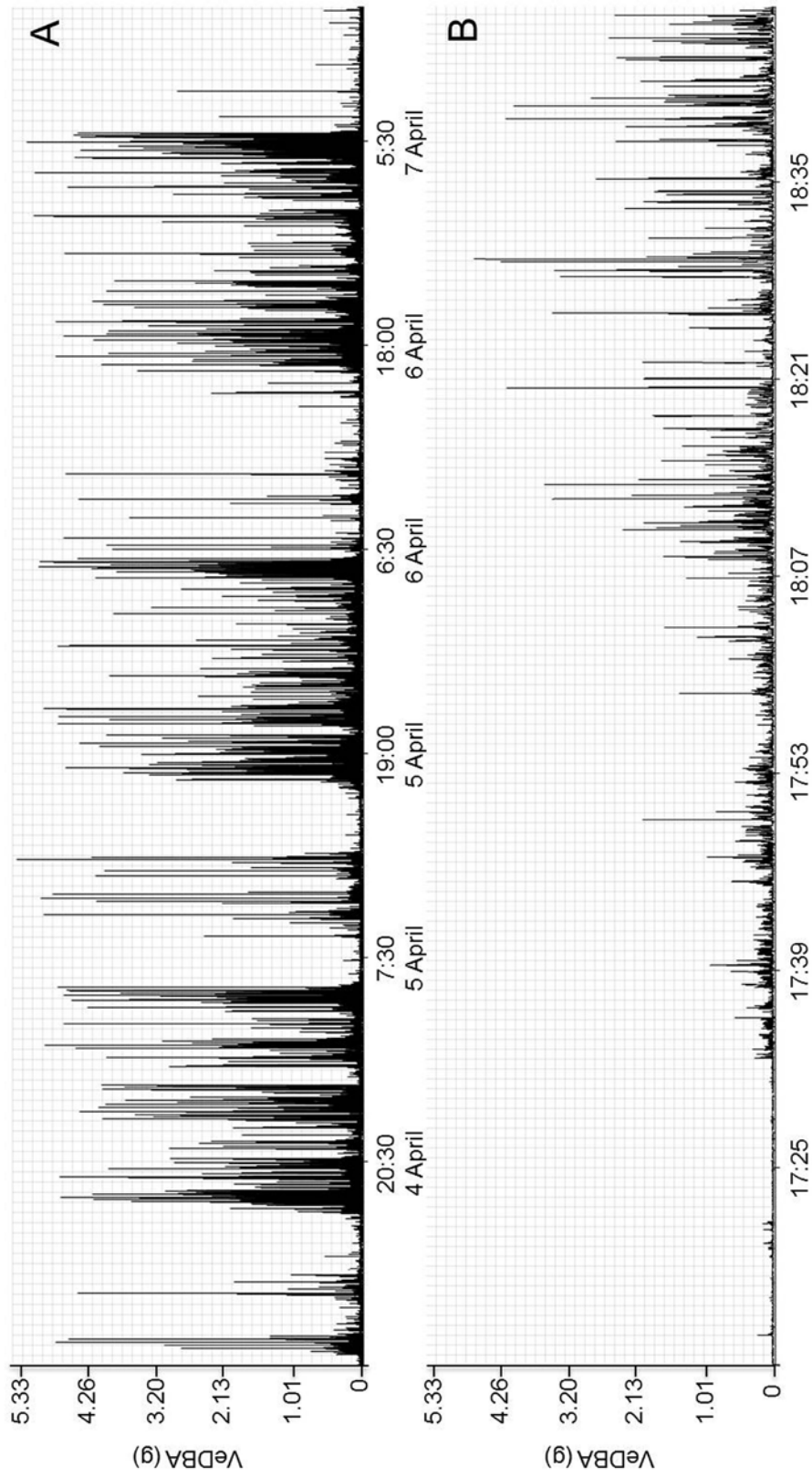


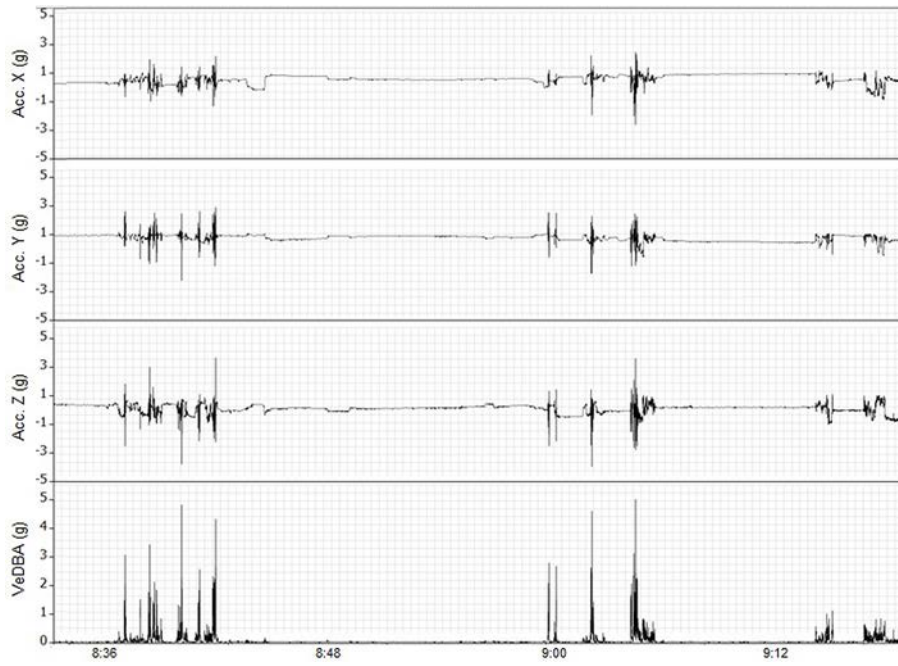
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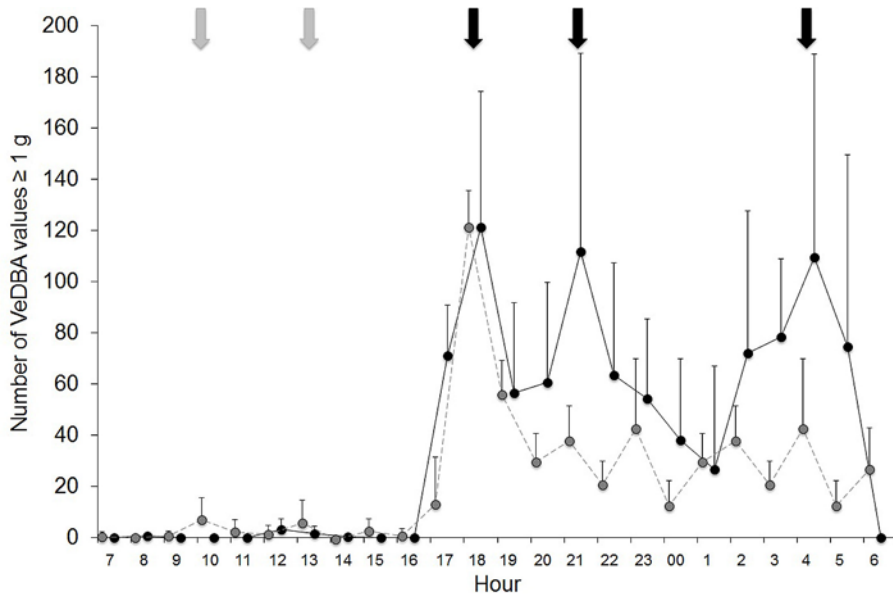
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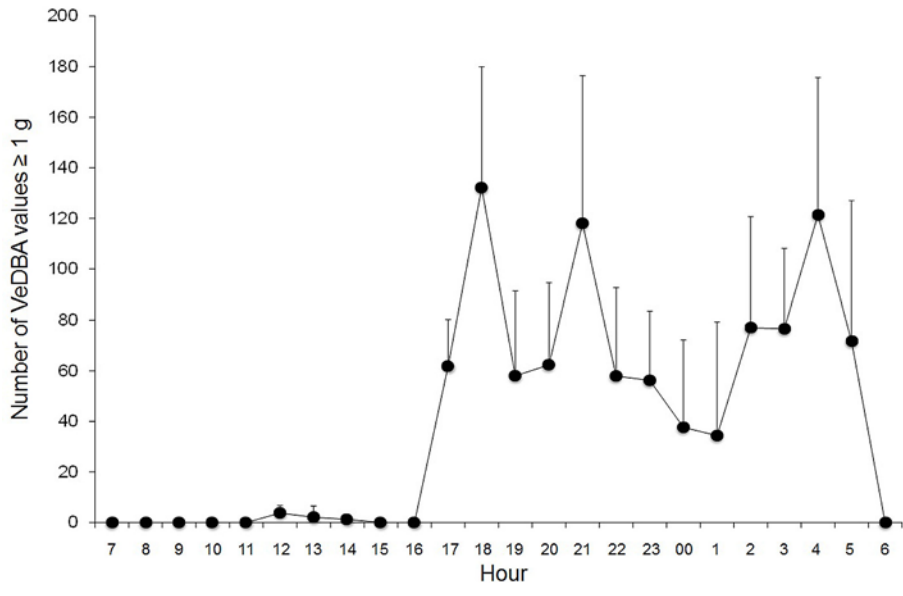
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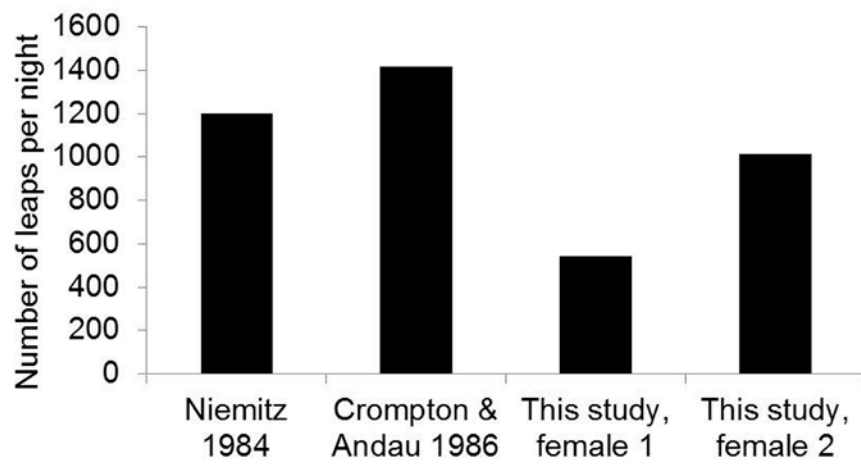
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