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## Supplementary Materials for

Summer Declines in Activity and Body Temperature Offer Polar Bears Limited Energy Savings

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21 **Materials and Methods**

22 Captures

23 Helicopter captures were performed in 2008 (August, October), 2009 (April–May,  
24 August, October), and 2010 (April–May). We searched for bears along the coast, on  
25 barrier islands, and up to 150 km offshore on sea ice (when present) between Barrow,  
26 Alaska (71°N, 157°W) and the Alaska-Yukon Territory border (69°N, 141°W). In  
27 October 2009 we also captured bears on sea ice between 70–79°N and 132–170°W, from  
28 the US Coast Guard Cutter *Polar Sea*. We immobilized bears with a mixture of tiletamine  
29 hydrochloride and zolazepam hydrochloride (Telazol; Warner-Lambert Co., Morris  
30 Plains, NJ) at estimated doses of 6 mg/kg of body mass, delivered in projectile syringes  
31 fired from a dart gun (33). During immobilization and processing we monitored rectal  
32 temperature and respiration rate. We weighed bears by suspending them in a net from a  
33 digital load scale hoisted from a tripod. Age was determined by counting cementum  
34 annuli in extracted vestigial premolars (34). Procedures for capturing, sampling, and  
35 instrumenting bears were approved by institutional animal care and use committees at the  
36 University of Wyoming and US Geological Survey (USGS) Alaska Science Center, and  
37 permitted by the US Fish and Wildlife Service (USFWS; Permit #MA690038). Use of  
38 trade names is for descriptive purposes only and does not imply endorsement by the US  
39 Government. This research was funded in part by the US Environmental Protection  
40 Agency (EPA) but because it was not reviewed by the EPA, no official endorsement  
41 should be inferred.

42

43 Location transmitters and collar data

44 Bears were instrumented in April–May (2009) and August (2008, 2009) and  
45 recaptured in October or the following spring. At recapture, instrumentation (see below)  
46 was retrieved and animals were released. Bears received one of four possible telemetry  
47 transmitters, either attached to collars (models TGW-3680 or TGW-4689H; Telonics,  
48 Mesa, AZ) or glued to the fur (SPOT or MK-10; Wildlife Computers, Redmond, WA).  
49 Collars automatically released 3–7 months after deployment and transmitters attached to  
50 fur detached during the molt or earlier. Collars recorded locations with Global  
51 Positioning System (GPS) technology (hourly; estimated error < 31 m; 35) or Doppler  
52 technology (typically every two days; estimated error: < 400 m, class 3; 401–1000 m,  
53 class 2; 1001–2499 m, class 1; 36). Glue-on transmitters were Doppler only. Limits to  
54 Argos satellite downloads yielded incomplete location datasets (36). Complete records  
55 were obtained from GPS collars retrieved from recaptured bears ( $n = 17$ ).

56 Both collar models were equipped with a temperature sensor (measurements every  
57 30 or 60 minutes, resolution 1°C). TGW-4689H also included an internal accelerometer,  
58 sensitive to motion in 3 planes, which recorded the total number of seconds of activity  
59 during the previous thirty minutes, every half hour. To 16 collars we attached an external  
60 accelerometer (Actiwatch; Mini-Mitter Respironics, Bend, OR) sensitive to motion in all  
61 planes, which recorded an acceleration score (a unitless index of acceleration intensity)  
62 every 2 minutes (37).

63

64 Temperature logger data

65 For recording abdominal temperature we used iButton (DS1922L, hourly  
66 measurements, resolution 0.0625°C; Maxim Integrated, San Jose, CA) loggers coated  
67 with impermeable paraffin wax and attached to a loop of sterile suture (0 gauge, non-  
68 dissolvable). Loggers were implanted into five bears fitted with location transmitters and  
69 activity loggers, and five bears with transmitters only. We shaved an area 10 × 6 cm on  
70 the ventral midline of each bear, approximately 80% of the distance from the xiphoid  
71 process to the umbilicus. We injected 1 mL of xylocaine into subcutaneous tissue  
72 throughout the shaved region, scrubbed the skin and surrounding fur with povidone-  
73 iodine. We then used sterile surgical tools to make a 7 cm incision through skin and  
74 subcutaneous adipose tissue to expose the linea alba. We made a 5 cm incision through  
75 the linea alba, pushed the logger through the incision and off to the side, and secured the  
76 logger to the underside of the linea alba, adjacent to the peritoneum, with a single stitch  
77 using the loop of suture. We closed the linea alba with interrupted single stitches, the  
78 adipose tissue with continuous stitches (00 gauge suture), and the skin with interrupted  
79 cruciate or mattress stitches. We used identical surgical procedures when recovering the  
80 loggers from recaptured bears.

81 Abdominal temperatures closely reflect intraperitoneal, or “core” temperature. In  
82 previous studies, measurements of abdominal temperature of polar bears (recorded from  
83 loggers in the same anatomical location as in our study; 38, 39) were nearly identical to  
84 intraperitoneal temperatures of polar bears in other studies (22, 40). The peritoneum itself  
85 tends to be very thin (< 1 mm in humans; 41) and offers little insulation.

86 To record rump temperature, we used Tidbit V2 loggers (measurements every 5 or  
87 10 minutes, resolution 0.02°C; Onset Computer Corporation, Bourne, MA) coated with  
88 wax and attached to a loop of sterile suture. Loggers were implanted in seven bears  
89 instrumented with transmitters and activity loggers. As described in Durner et al. (6), we  
90 made an incision through the skin approximately 15 cm to one side and 5 cm ventral of  
91 the base of the bear’s tail. We separated the subcutaneous adipose tissue and secured the  
92 logger to the surface of the gluteus maximus muscle using the loop of suture. Loggers  
93 were recovered using identical procedures.

94 After abdominal and rump loggers were retrieved, we placed them in a temperature-  
95 controlled chamber (CL-740A; Omega Engineering Inc., Stamford, CT) and created 4–5  
96 point calibration curves for the range of temperatures recorded from each bear. Next, we  
97 left loggers in an environmental chamber (Convicon, Pembina, ND) held at a steady  
98 temperature for eight weeks and confirmed that measurements did not drift over time.

#### 100 Calculating smoothed body temperatures

101 We created smoothed values of abdominal and rump temperatures using seasonal  
102 trend decomposition with the “stl” (seasonal trend loess) command in the base statistical  
103 package in Program R (54). The seasonal window was periodic and the loess window  
104 was 25% of the average sample size of bear records extending from May to October.

#### 106 Comparing data from bears on shore and ice

107 To ensure capture effects and incision healing did not influence results, for all  
108 analyses we censored all data collected within 120 hours (5 days) after capture,  
109 encompassing the period during which polar bears return to regular movement patterns  
110 (42, 43), and within 1 hour prior to recapture. Data of movement rate, acceleration scores,

111 collar temperature, rump temperature, and body mass for bear 20741, which engaged in a  
112 nine-day swim in August–September 2008, were previously reported in a companion  
113 study (30). Such a reported long-distance swim is unique (13) thus we excluded these  
114 data from calculations and statistical tests of mean movement rate and activity (Fig. 2 and  
115 table S1). However, we used these data in models to identify the variables that predicted  
116 acceleration scores (table S3) and to evaluate the relationship between acceleration scores  
117 and body temperatures (table S5).

118 To assess activity trends over time, we calculated a single activity variable: the  
119 proportion of time spent active. To derive this proportion from measurements of the  
120 number of seconds of activity per 30 minutes, we divided the count by 1800. For  
121 acceleration scores, we first converted the scores to the number of seconds of activity per  
122 30 minutes. To achieve this conversion, we calculated a mean acceleration score in half-  
123 hour blocks. Next, we pooled data for seven bears which had both types of activity  
124 measurements and regressed the number of seconds of activity during the previous half  
125 hour against the mean acceleration score for that half-hour (total pooled measurements,  $n$   
126 = 27,579). The relationship appeared linear for small values but increasingly non-linear at  
127 high values, thus we used automated segmented regression (44) and identified a mean  
128 acceleration score of 1128 as the breakpoint between linear segments (fig. S4A). For  
129 scores below this value, the two types of activity data were related by this equation:

130

131  $N$  = number of seconds of activity over the previous half-hour

132  $M$  = mean acceleration score over the previous half-hour

133  $N = (M \times 0.838) + 38.645$

134

135 All monthly means of 30-minute mean acceleration scores were  $< 1128$ , thus we  
136 used the above equation to convert these data into monthly means of the number of  
137 seconds of activity over the previous half-hour. For bears with both types of activity data,  
138 the monthly means of the proportion of time spent active based on measured seconds of  
139 activity were similar to the monthly means of the proportion of time spent active based  
140 on acceleration scores converted to seconds of activity ( $n = 30$ , pooled data from seven  
141 bears; fig. S4B). This indicates the converted data adequately represents the time spent  
142 active.

143 We calculated monthly means of the time spent active (after pooling measured and  
144 converted data), movement rate, and smoothed abdominal temperature for each bear, for  
145 each month in which it had  $\geq 96$  hours of measurements. Each monthly mean was  
146 categorized as “Shore” or “Ice” based on GPS and Doppler (class 2 and 3) locations.  
147 Three bears with location and activity data moved between habitats in July–October, and  
148 we calculated a monthly mean for both Shore and Ice accordingly. We compared Shore  
149 and Ice monthly means where  $n \geq 3$  for each group, using the Welch t-test (45).

150

#### 151 Modeling influence of environmental variables on activity

152 We used ARIMA (autoregressive, integrated, moving average) approaches (“arima”  
153 command, base statistical package in Program R; 46) to model the influence of three  
154 environmental predictor variables on both types of activity data: daily mean of the  
155 number of seconds of activity per half-hour, and daily mean of the acceleration scores.  
156 Individual models were built for each bear.

157 The first model (ARIMA structure [1,0,1]) was built with the predictor “Whale”  
158 (only applied to bear locations on shore), a variable between 0–1, describing the daily  
159 proportion of hourly GPS locations < 500 m from the site of a whale carcass, ≤ 60 days  
160 after the landing of a whale at that site (47, 48). Although bears may scavenge on these  
161 carcasses year-round (49), they likely consume the majority of tissue shortly after whales  
162 are landed (50).

163 In the second model (ARIMA structure [1,0,1]) we used the predictor “Air  
164 Temperature” (only applied to locations on shore), a daily mean of hourly air  
165 temperatures recorded < 200 km from each bear location at Alaskan weather stations in  
166 Kaktovik, Deadhorse, or Barrow (www.weatherspark.com).

167 In the third model (ARIMA structure [1,0,1]) we used the predictor “Shelf” (only  
168 applied to locations on the sea ice), a binary variable indicating whether the daily mean  
169 bathymetry value of bear locations was ≥ 300 m. Shelf edges are abrupt in this region of  
170 the Arctic and habitat studies indicate that bears select for shallow waters (51, 52). Water  
171 depths at bear locations were assigned based on the International Chart of the Arctic  
172 Ocean 3.0 (53).

173

#### 174 Correlating activity and body temperatures

175 Abdominal temperature was recorded hourly and rump temperature every 5 or 10  
176 minutes, but neither was synchronized with the hourly measurement of other variables  
177 (e.g., seconds of activity in the previous half-hour). Thus, for correlating temperatures  
178 with activity, we used linear interpolation to estimate body temperatures at the exact time  
179 activity was measured.

180 We evaluated the correlation between activity and body temperatures using  
181 seemingly-unrelated-time-series models, with commands in the “dlm” (dynamic linear  
182 models) package in Program R (55, 56). Models were developed, by bear, with two  
183 activity variables matched with interpolated values of the two body temperature variables  
184 (abdominal and rump temperature) at an hourly scale. Activity included: a) seconds of  
185 activity in the previous half hour, and b) mean acceleration score over the previous half  
186 hour.

187 For these models we used adjusted abdominal and rump temperatures, which only  
188 included interpolated values within 4 SD of the smoothed temperature; this excluded  
189 most outliers of cold temperatures recorded during swimming events (described below).  
190 The initial models correlating both measures of activity to rump temperature were  
191 unstable and could not provide inferences, likely because rump temperature declined  
192 substantially during some periods of inactivity. Thus, we censored the data to only  
193 include rump temperature measured when the mean acceleration score during the  
194 previous half-hour was ≥ 30. We used mean acceleration scores because all bears with  
195 rump temperature measurements had these data, but not all had measurements of number  
196 of seconds of activity. We selected this cutoff (≥ 30) after inspecting the data and  
197 observing periods of several hours where scores were mostly 0, indicating motionless  
198 bears, but which were interrupted by occasional large acceleration scores (e.g. up to 100),  
199 likely representing resting interspersed with small movements such as grooming. We  
200 reasoned that half-hour means of scores recorded during resting were unlikely to exceed  
201 30. Pooled half-hour means of acceleration scores from all bears ranged from 0–4933,  
202 with a grand mean (95% CI) across all bears of  $454 \pm 6$ .

203

204 Cold abdominal and rump temperatures during inactivity and swimming

205 We identified periods of inactivity as  $\geq 2$  hours with mean hourly movement rate  $<$   
206 0.01 m/s and mean acceleration scores  $< 30$ , and during these periods we counted the  
207 instances of hourly, interpolated abdominal and rump temperatures that were  $< 35.0^{\circ}\text{C}$ .  
208 We used interpolated temperature measurements to match the timing of recording of  
209 movement rates and activity.

210 We identified periods of swimming from locations in open water, or time periods  
211 occurring between sequential locations moving to or from an island. All identified  
212 swimming events occurred near shore during summer, reducing the potential for  
213 occurrence of ice floes large enough for bears to walk on but small enough to escape  
214 satellite detection. We verified that during each swim, surrounding water had no sea ice  
215 and collar temperatures were between  $-3^{\circ}\text{C}$  and  $+12^{\circ}\text{C}$ , the range of feasible temperatures  
216 of surface water in this region of the Arctic. If a swim began between hourly locations,  
217 we evaluated acceleration scores during that period. The start of a swim was marked by  
218 elevated, steady scores with little variation. The end of a swim was typically marked by a  
219 sudden increase in variability of scores, likely reflecting a bear shaking or rolling on the  
220 ground or ice to shed water from its fur. During periods of swimming, we counted the  
221 instances of abdominal temperature (measured hourly) and rump temperature (measured  
222 every 5 or 10 minutes) that were  $< 35.0^{\circ}\text{C}$ . Many potential swimming events were  
223 discarded because they did not meet all criteria. Similarly, we did not attempt to identify  
224 swimming events for bears on the sea ice because ice data were too coarse to determine  
225 whether bears were walking on ice or swimming between floes.

226

227 Evaluating the cooling rate of abdominal loggers

228 The maximum passive cooling rate of tissue occurs after death, and equations  
229 describing cooling of human cadavers are used to estimate time of death in forensic  
230 sciences (57). These equations have been modified to reflect cooling after death of  
231 marine mammal carcasses, using experimental data collected from intraperitoneal loggers  
232 implanted into California sea lion (*Zalophus californianus*) carcasses immersed into  
233 temperature-controlled water (58). Agreement between modeled and experimental data  
234 indicates that the primary influences on cooling rate are carcass size and water  
235 temperature (58), suggesting that these equations can reasonably be applied to polar  
236 bears. We calculated a theoretical cooling curve for bear 21150, under the pretense that it  
237 died on October 3<sup>rd</sup>, using these assumptions: bear temperature at time of death of  
238  $38.0^{\circ}\text{C}$ , matching the actual abdominal temperature recorded at that time; body mass of  
239 174 kg, based on a linear rate of change between August 10<sup>th</sup> (when this bear was  
240 measured at 123 kg) and October 18<sup>th</sup> (measured at 188 kg); and immersion in moving  
241 water of  $4^{\circ}\text{C}$  (typical coastal Arctic sea water in summer). We compare the theoretical  
242 curve to observed data from the live bear in Fig. 4.

243

244 **Supplementary Text**

245 Estimating heat loss and skin temperature of polar bears while swimming

246 We calculated surface area of a bear as  $0.09/(\text{body mass in kg})^{0.67}$  and assumed 98%  
247 of that area (everything but the head) was submerged (59). Heat losses from the head

248 (radiative, convective, respiratory) were ignored, as they have been shown to be minor  
249 (22, 59) in comparison to conductive heat loss in water. We assumed the water  
250 temperature was 4°C, the bear was swimming at the typical rate of 0.56 m/s (13), and  
251 conductance across the skin and fur was 128.6 W/m<sup>2</sup>/°C. The latter is based on an  
252 increase in conductance across polar bear skin and fur (summer pelage) of 3.4×10<sup>-4</sup>  
253 calories/cm<sup>2</sup>/second/°C for every 0.1 m/s increase in water speed (60). In calculations we  
254 varied the skin surface temperature from 5°C to 30°C.

255 Polar bears swim using alternate pectoral paddling, which is more efficient than  
256 quadrupedal paddling (61). Polar bear metabolic rates while swimming have not been  
257 measured, but may be similar to their metabolic rates while walking. For ferrets (*Mustela*  
258 *putorius furo*), the only other mammal known to predominantly use alternate pectoral  
259 paddling, rates differ by < 15% between swimming and walking (61, 62). In a previous  
260 captive experiment, 125-kg and 155-kg polar bears had metabolic rates of 125 W and 175  
261 W at rest, and 528 W and 891 W when walking at 1.0 m/s (63). Under the assumption  
262 that metabolic rates of walking and swimming are identical, linear interpolation yields  
263 rates of 351 W and 576 W for travel at 0.56 m/s. The mean mass-specific cost at this  
264 swimming speed is thus 2.79 W/kg, and a 300-kg bear would then have a metabolic rate  
265 of 837 W. Under these assumptions, bears require skin surface temperatures < 6°C to  
266 avoid conductive heat losses that exceed metabolic heat production (fig. S5).

267

#### 268 Body temperature during pregnancy and winter hibernation

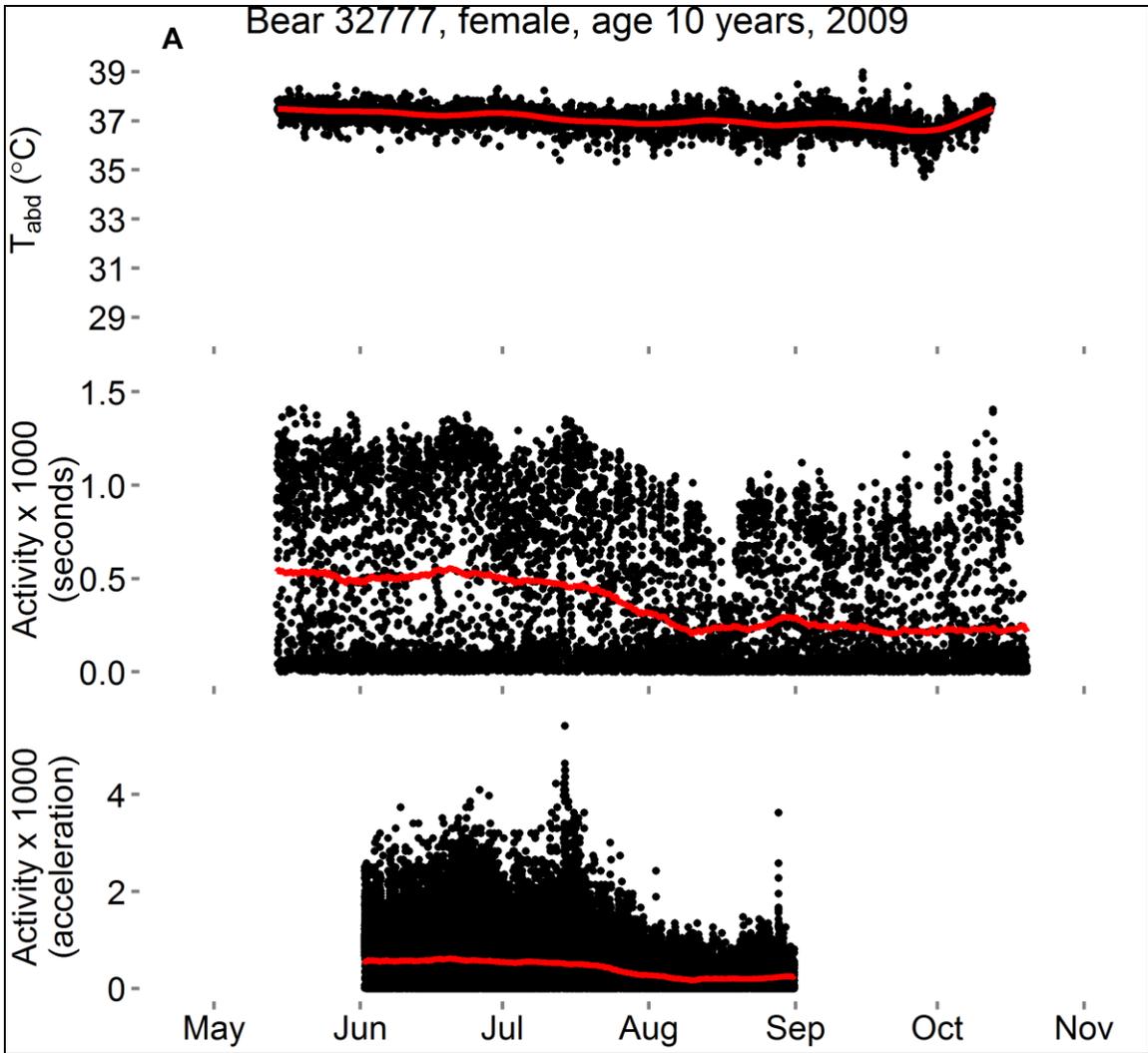
269 Bear 20529 was implanted with an abdominal temperature logger on August 11<sup>th</sup>  
270 2009 and poor weather prevented her recapture in October. She was recaptured with two  
271 cubs-of-the-year on April 5<sup>th</sup> 2010. Her logger ceased recording when the memory filled  
272 on January 19<sup>th</sup>. Her temperature profile is shown in figures S1 and S2B. Similar to other  
273 bears, her temperature declined through August until mid-September. It then abruptly  
274 increased in early October, suggesting blastocyst implantation and the need for high and  
275 stable temperatures during fetal development (26, 64). She likely entered a maternity den  
276 on November 3<sup>rd</sup>, as indicated by Doppler locations clustered in the same area from  
277 November 3<sup>rd</sup> to 23<sup>rd</sup>, when the location transmitter failed. Her temperature declined  
278 slightly through November then fell abruptly around November 28<sup>th</sup>, suggesting a 55–60  
279 day gestation with a small and progressive reduction in core temperature, identical to  
280 brown bears (26). From December 6<sup>th</sup> to January 19<sup>th</sup>, her mean (± 95% CI) hourly  
281 temperature was 35.0°C (± 0.02°C).

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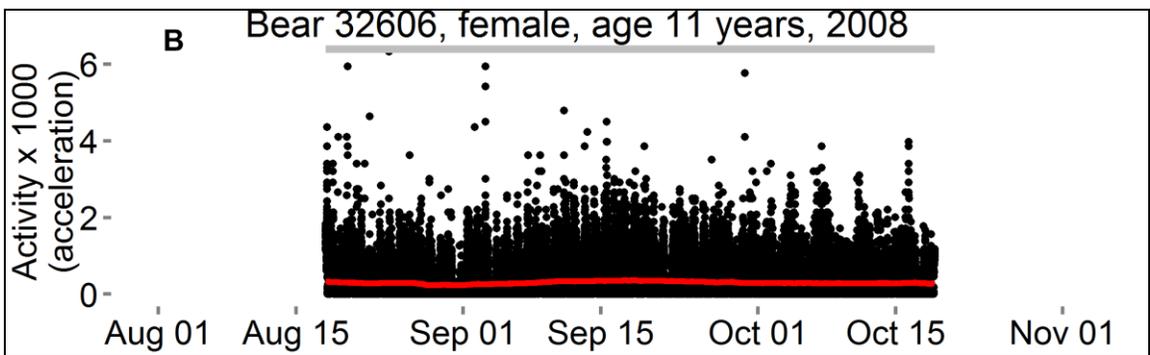
#### 283 Describing greater temperature swings in the rump than in the abdomen

284 The mean of the maximum hourly increase recorded by rump loggers (+11.8°C;  
285 from  $n = 7$  bears) was greater than the mean of the maximum hourly increase recorded by  
286 abdominal loggers (+5.0°C;  $n = 10$ ), based on a Mann-Whitney Rank Sum test ( $p =$   
287 0.003,  $U = 4.0$ ,  $T = 94.0$ ). Similarly, the mean hourly decrease was also larger for rump ( $-$   
288 10.4°C;  $n = 7$ ) than abdominal (-5.0°C;  $n = 10$ ) loggers (t-test,  $p = 0.006$ ,  $t = -3.23$ ,  $df =$   
289 15).

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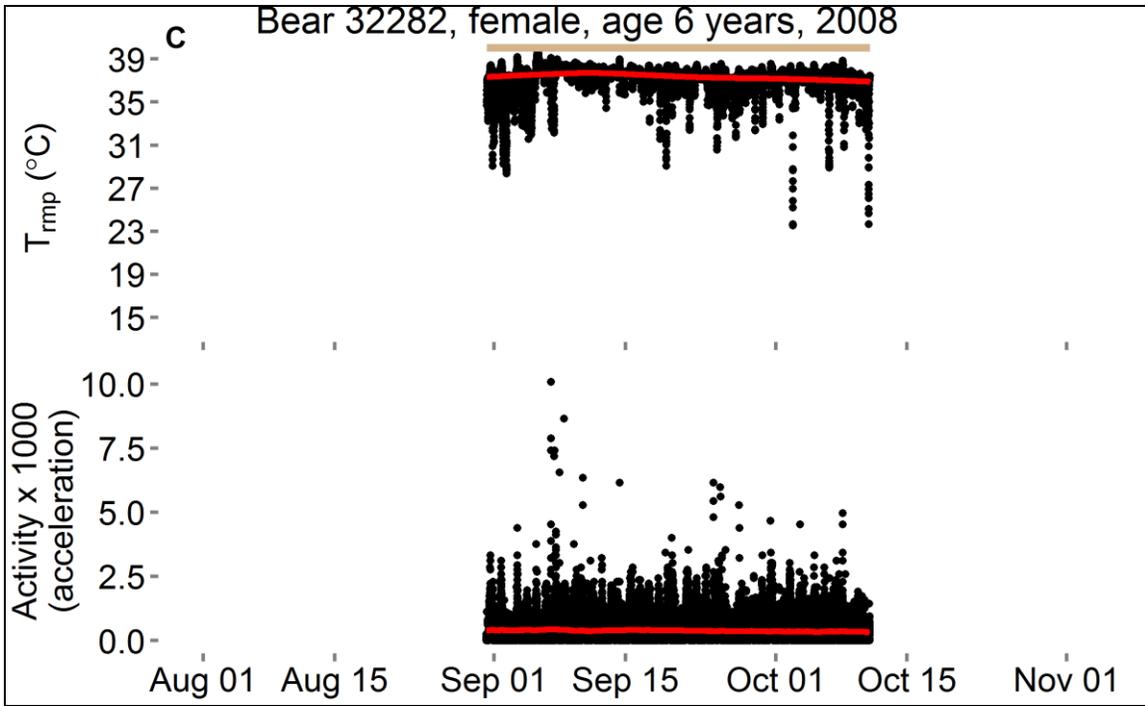


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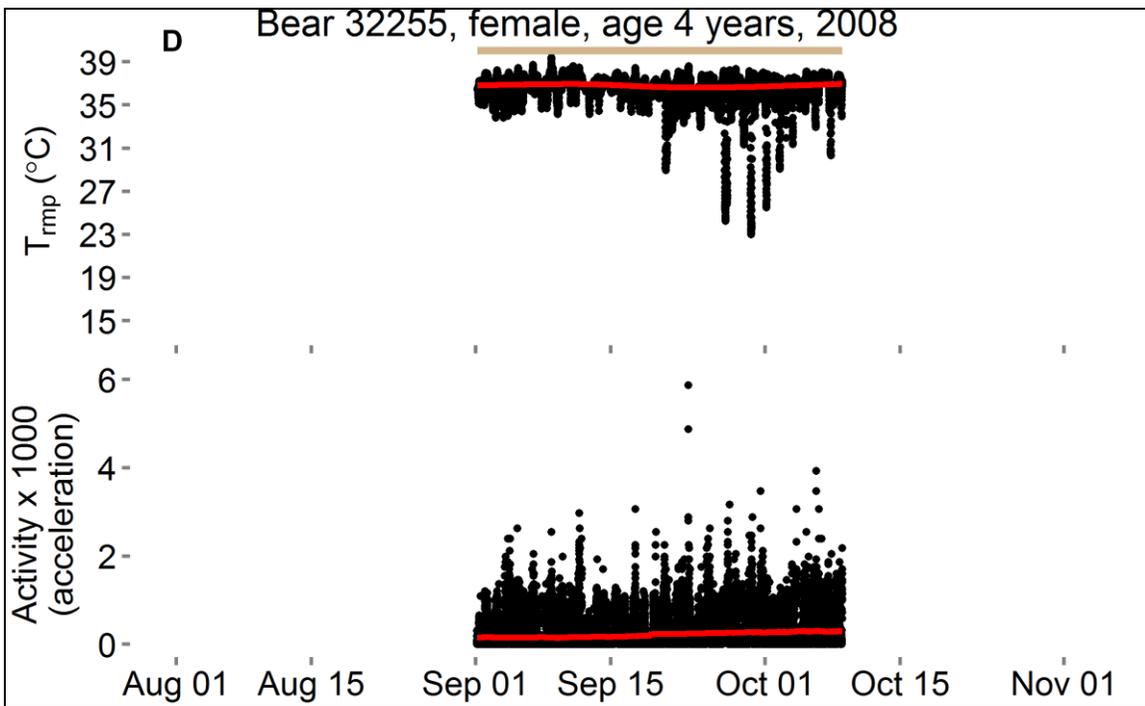


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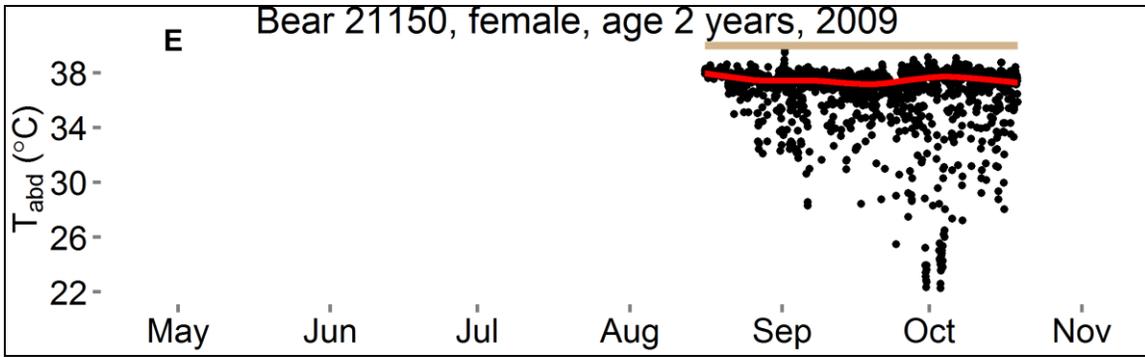
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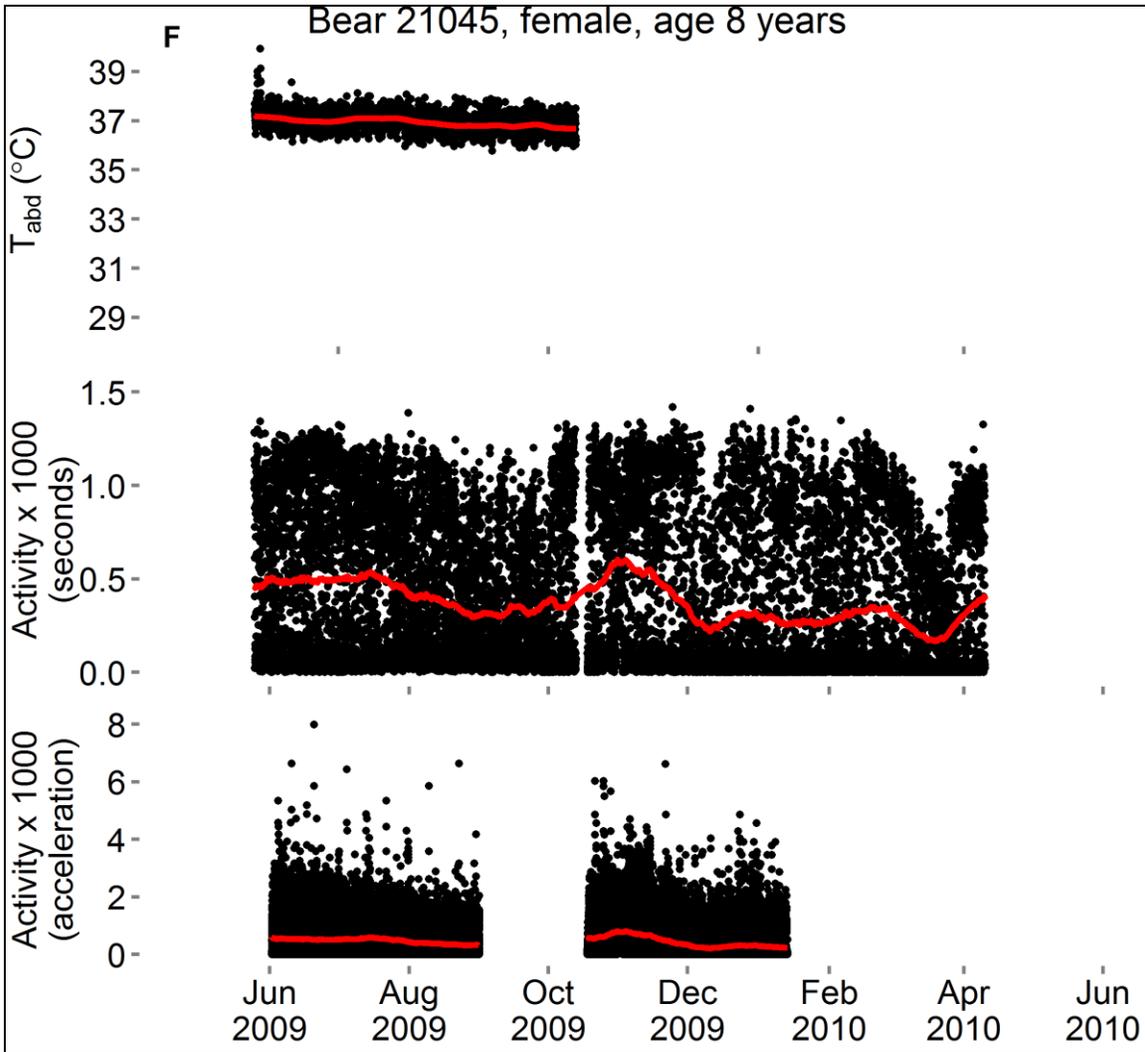
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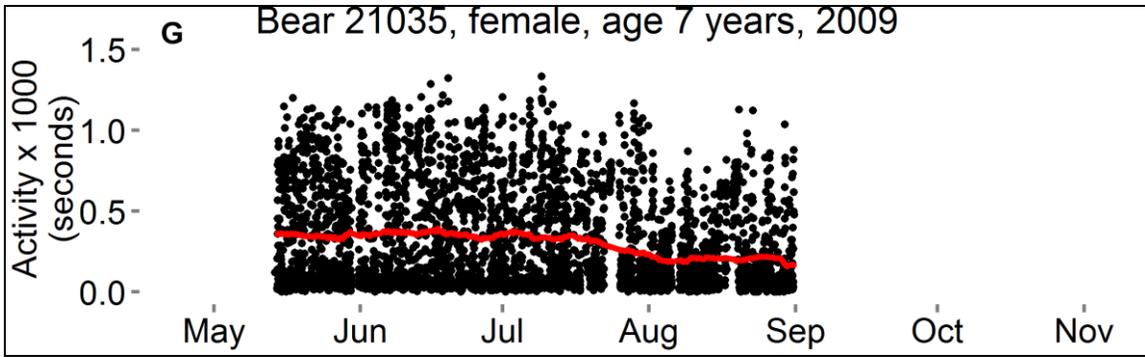
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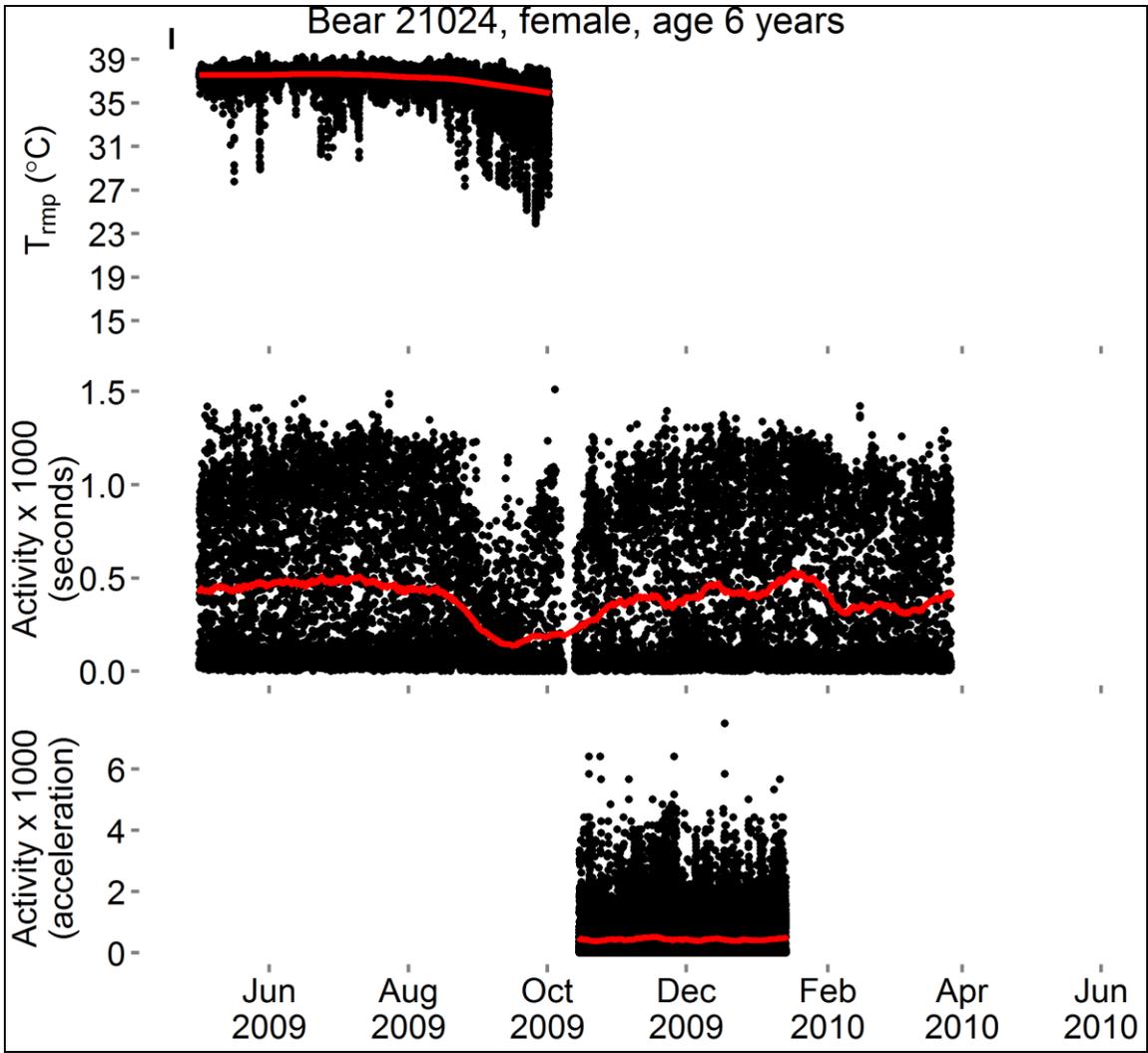


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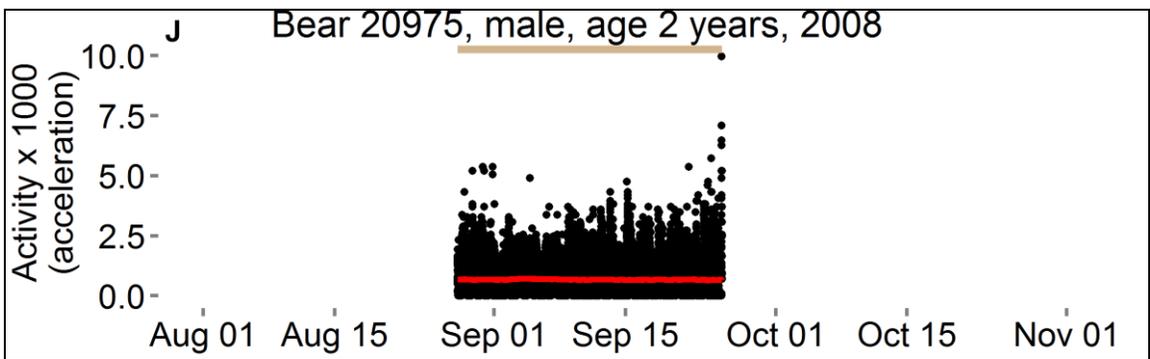


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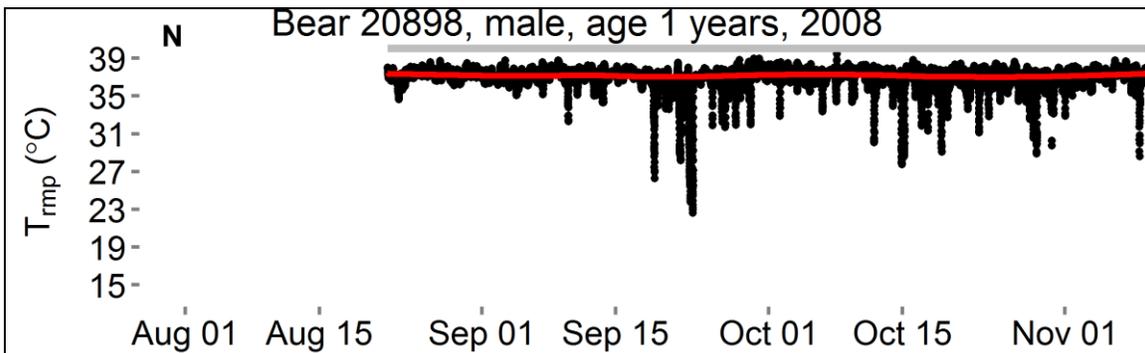
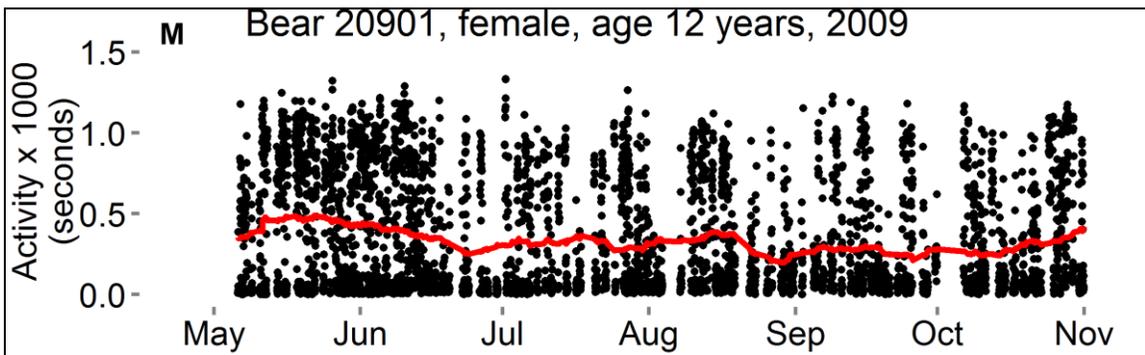
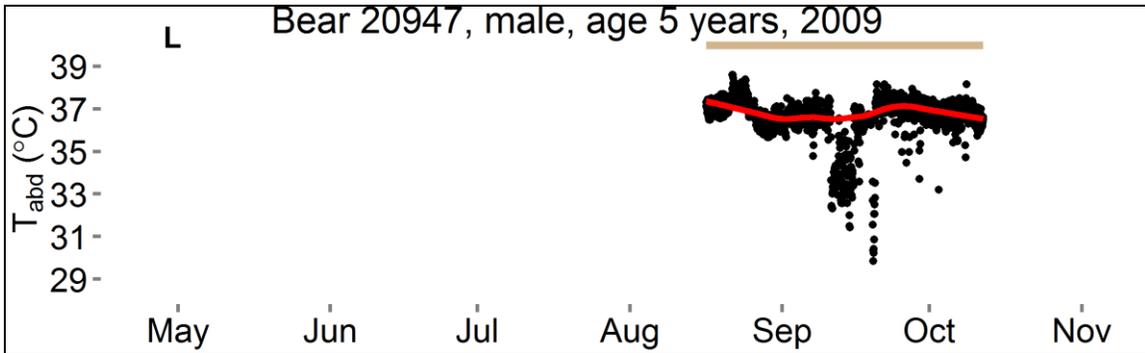
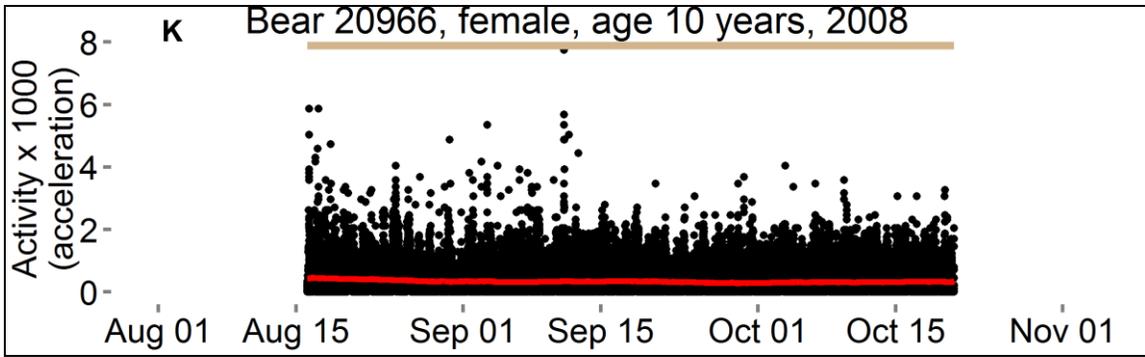


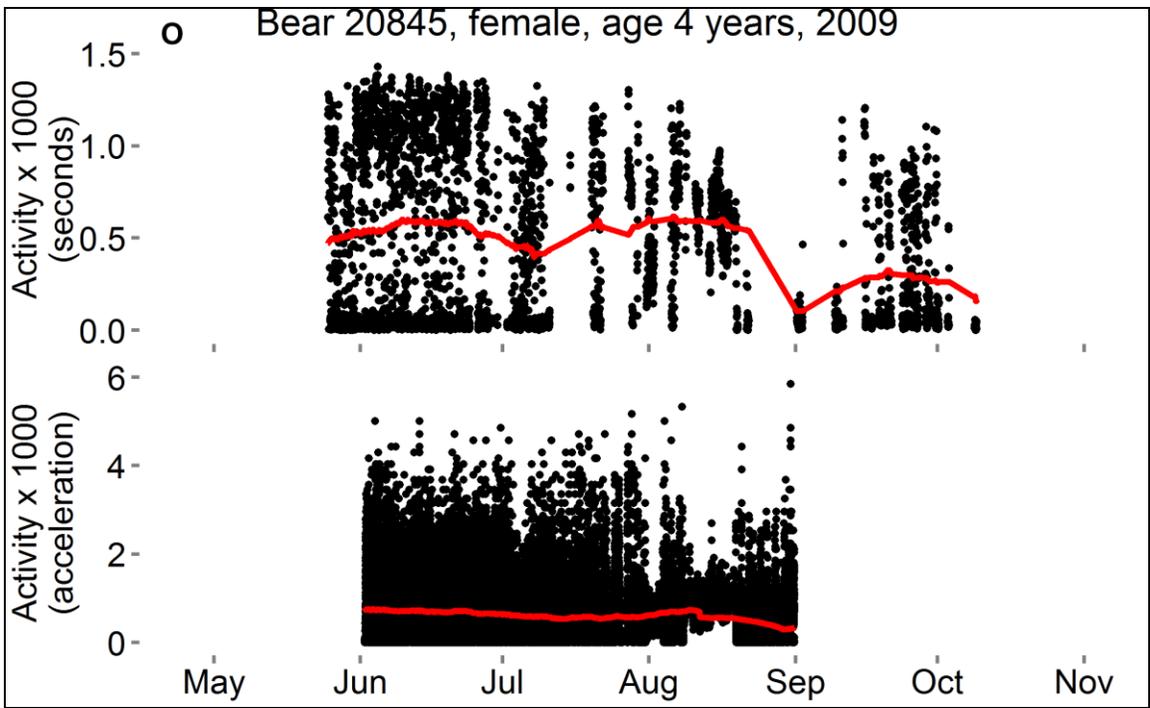


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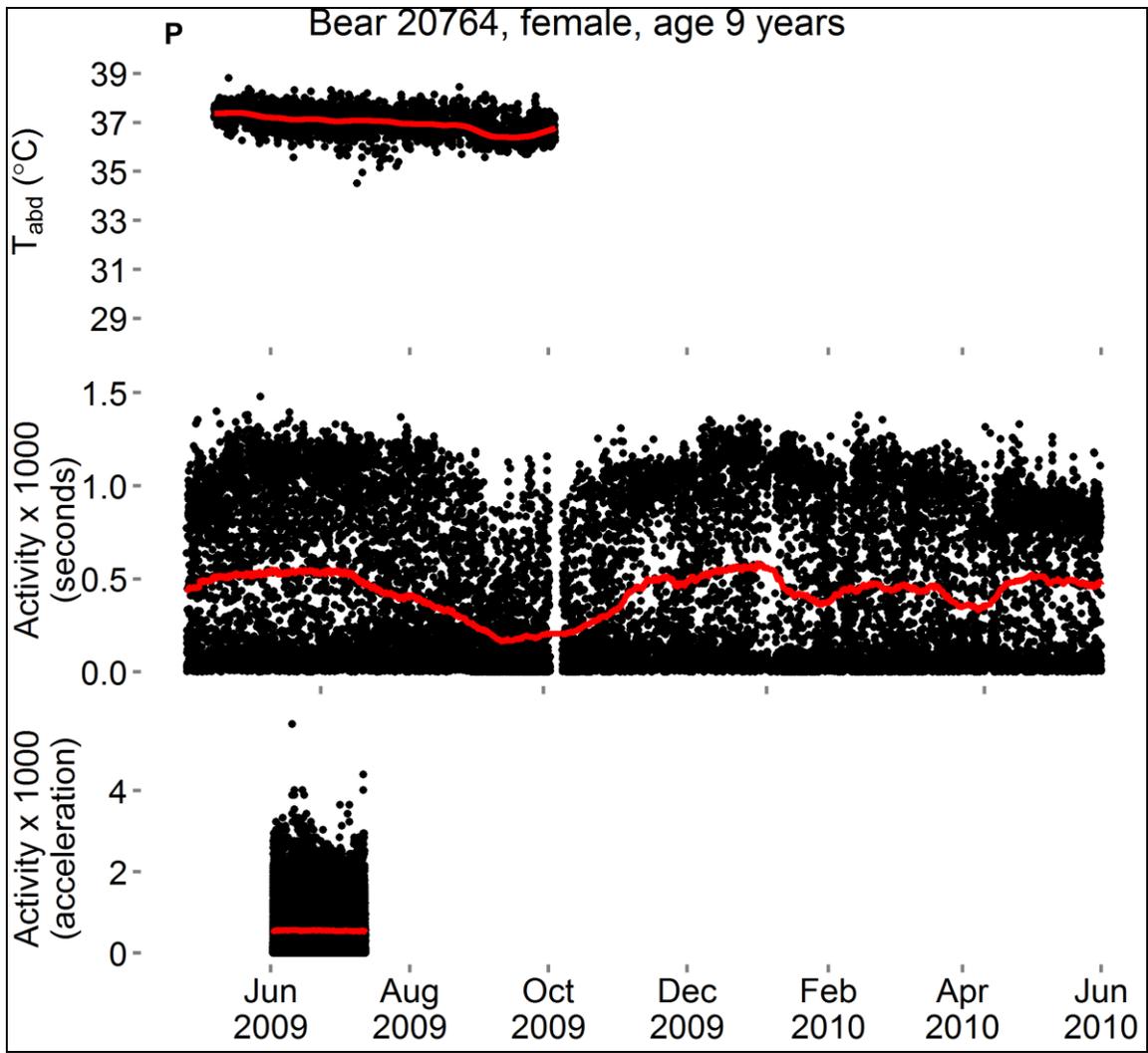


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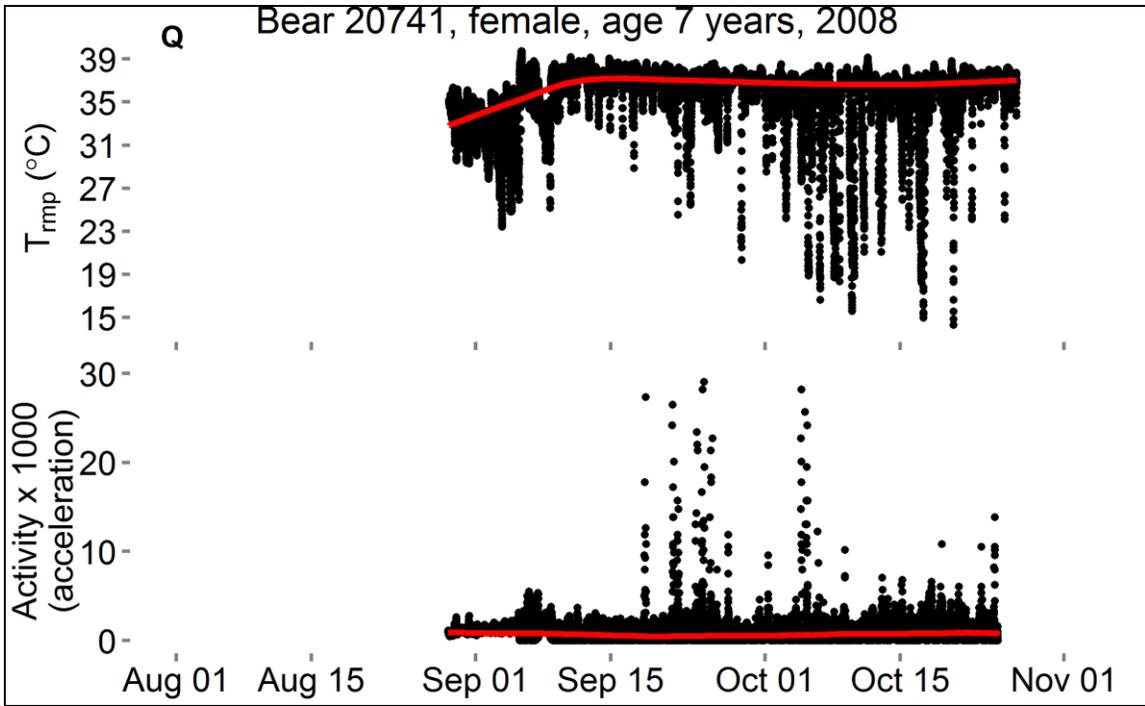


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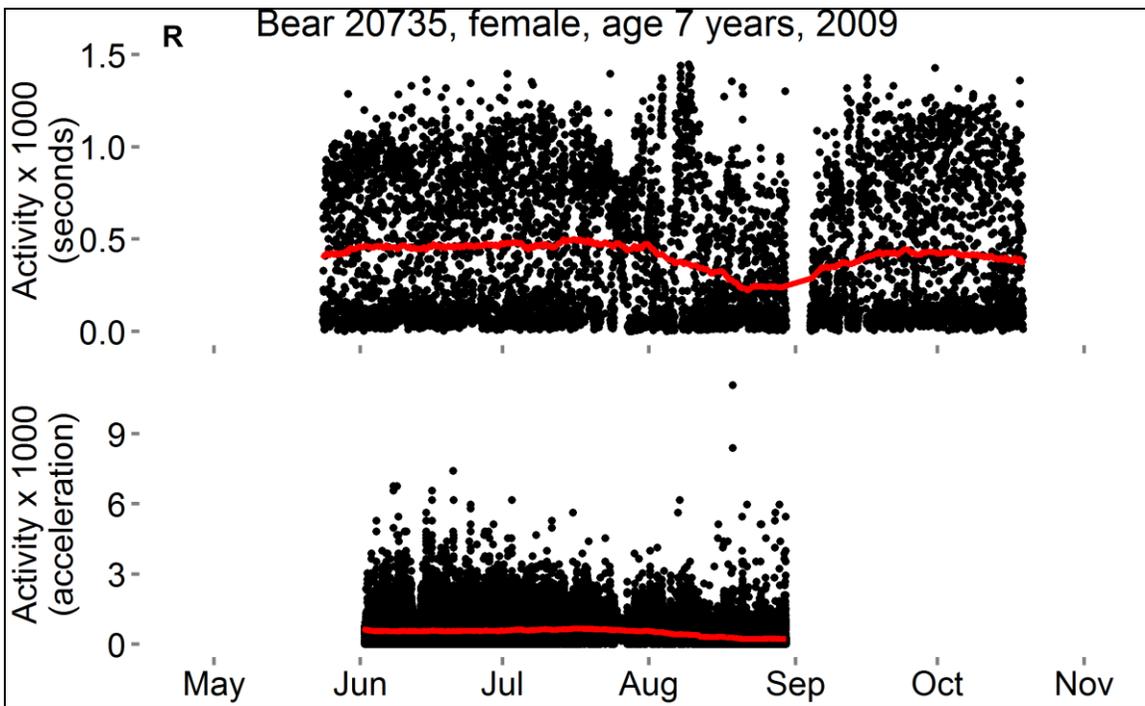


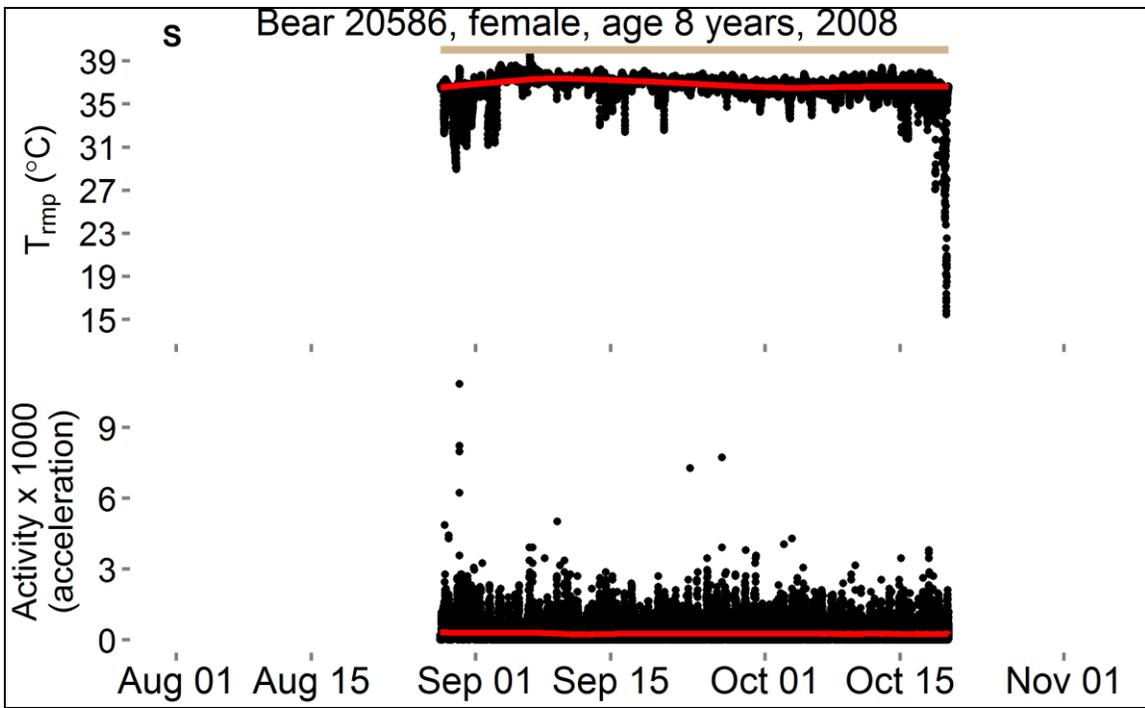
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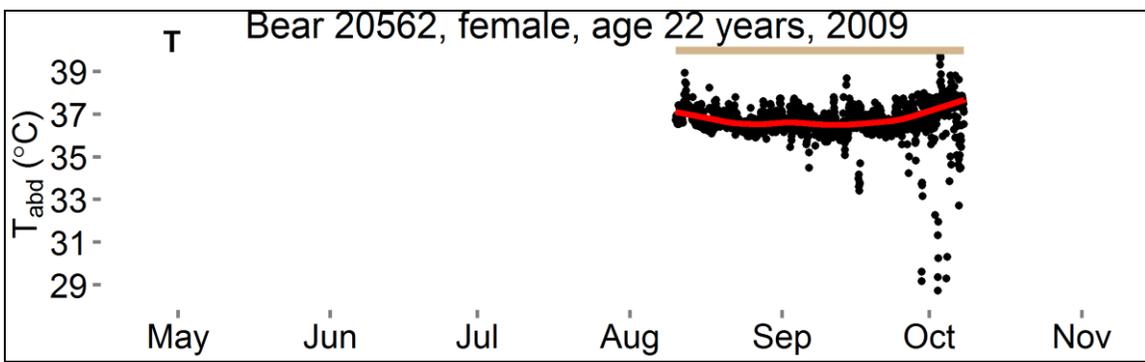


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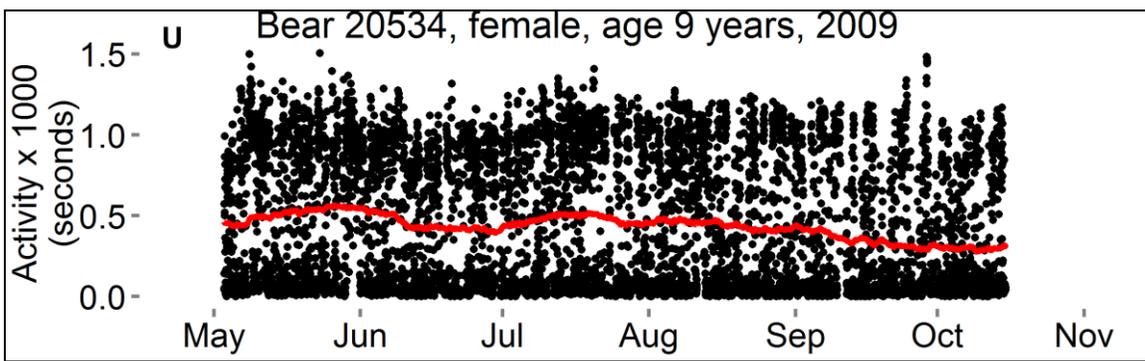




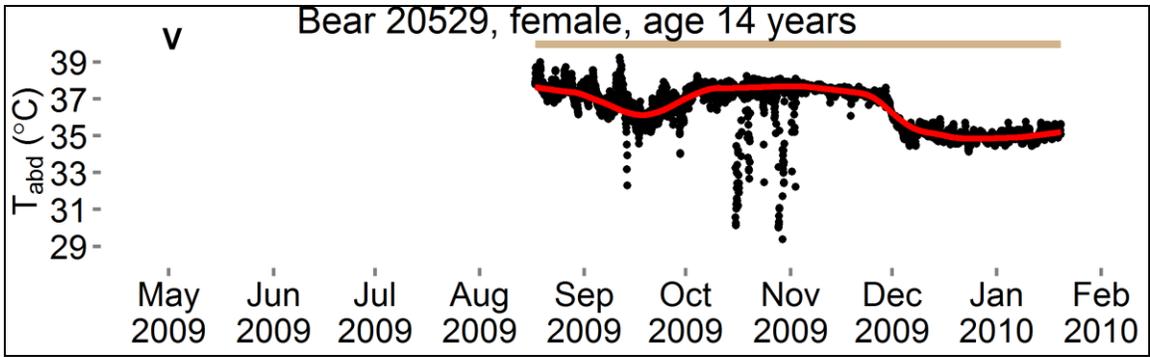
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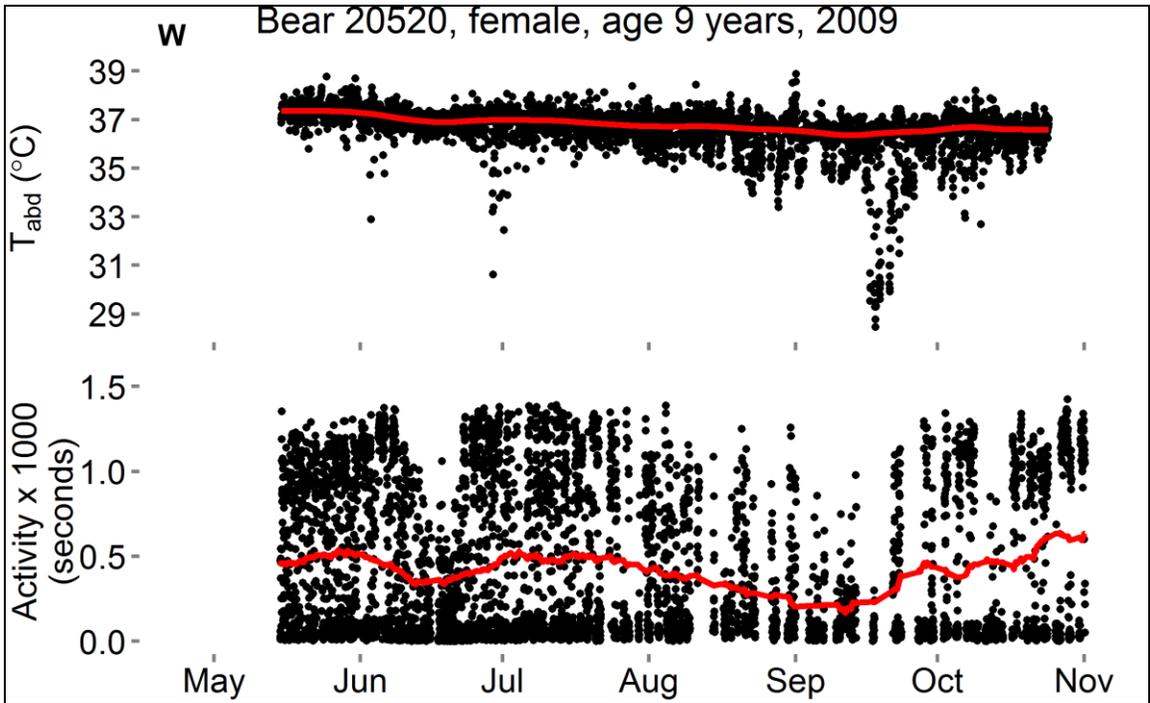
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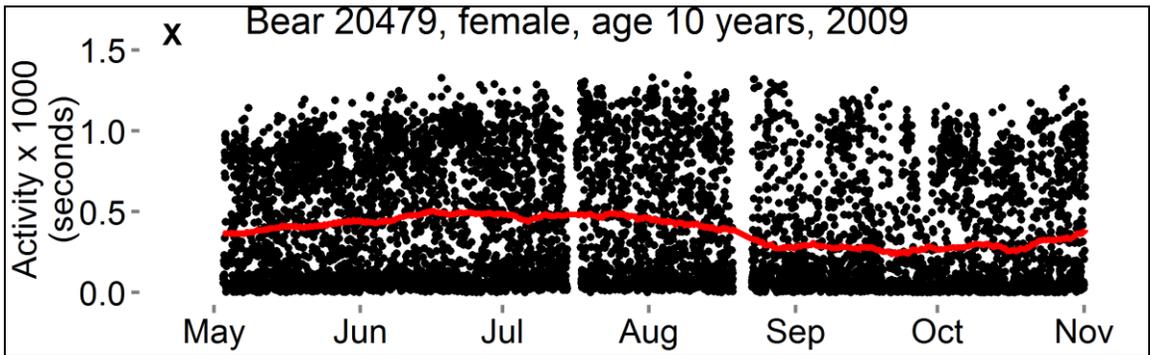
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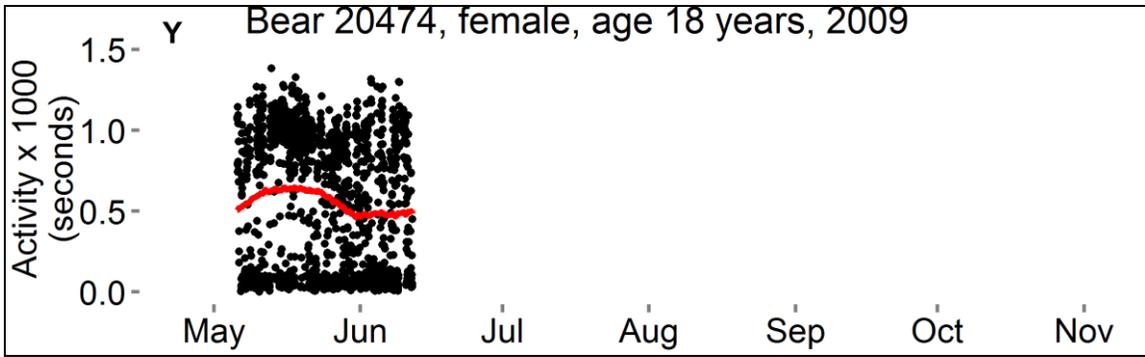


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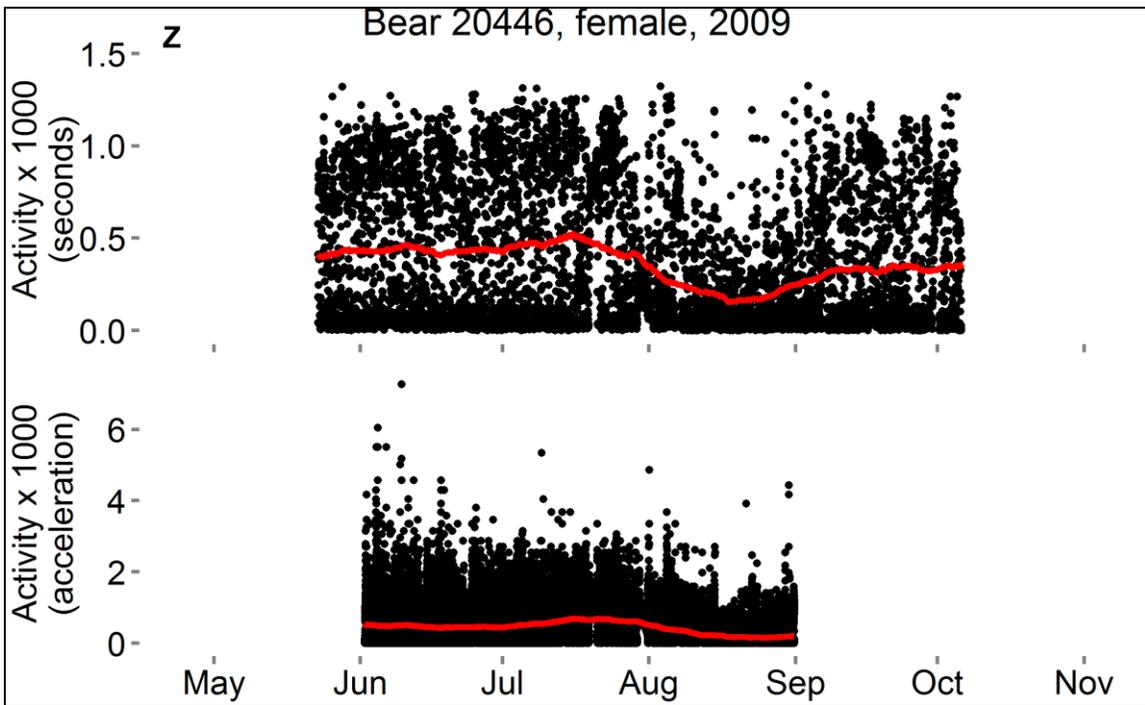


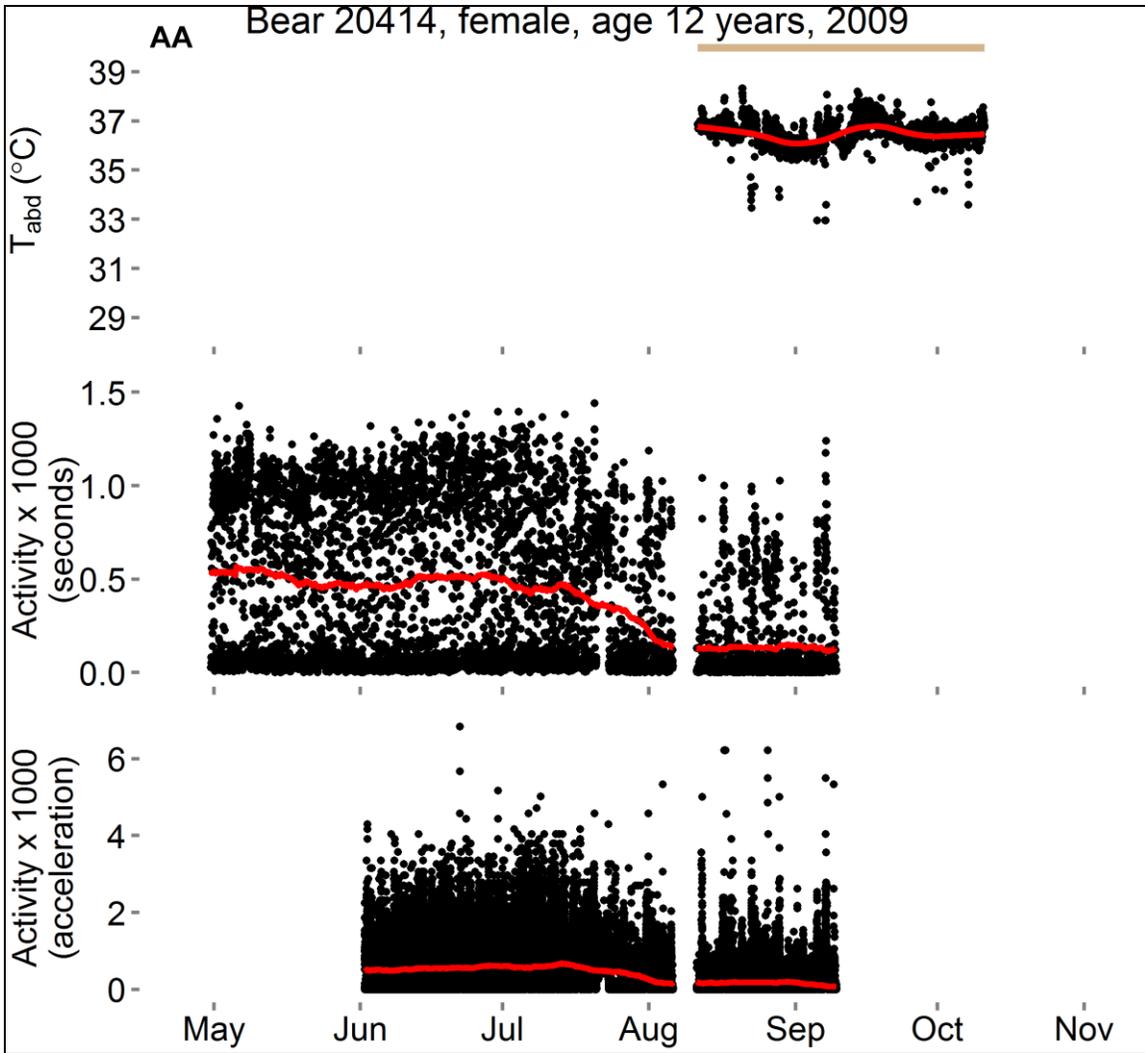
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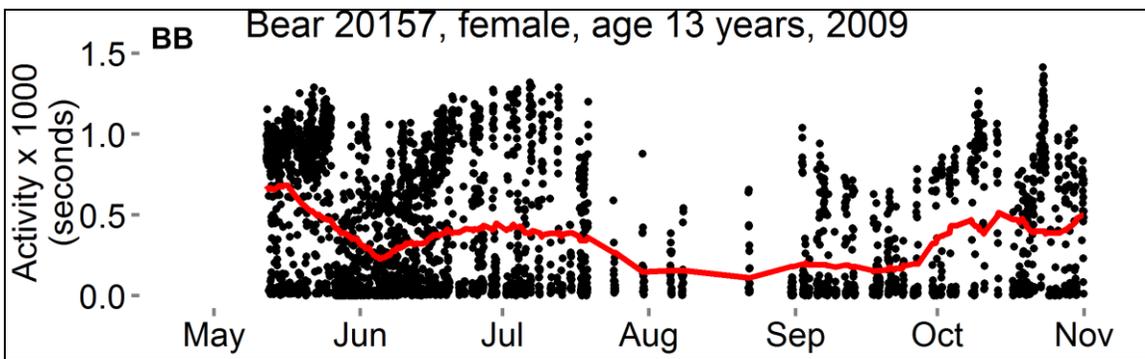


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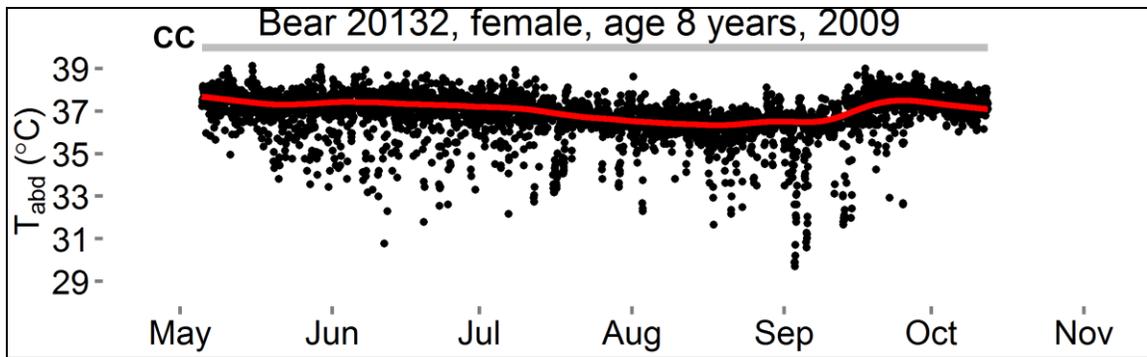


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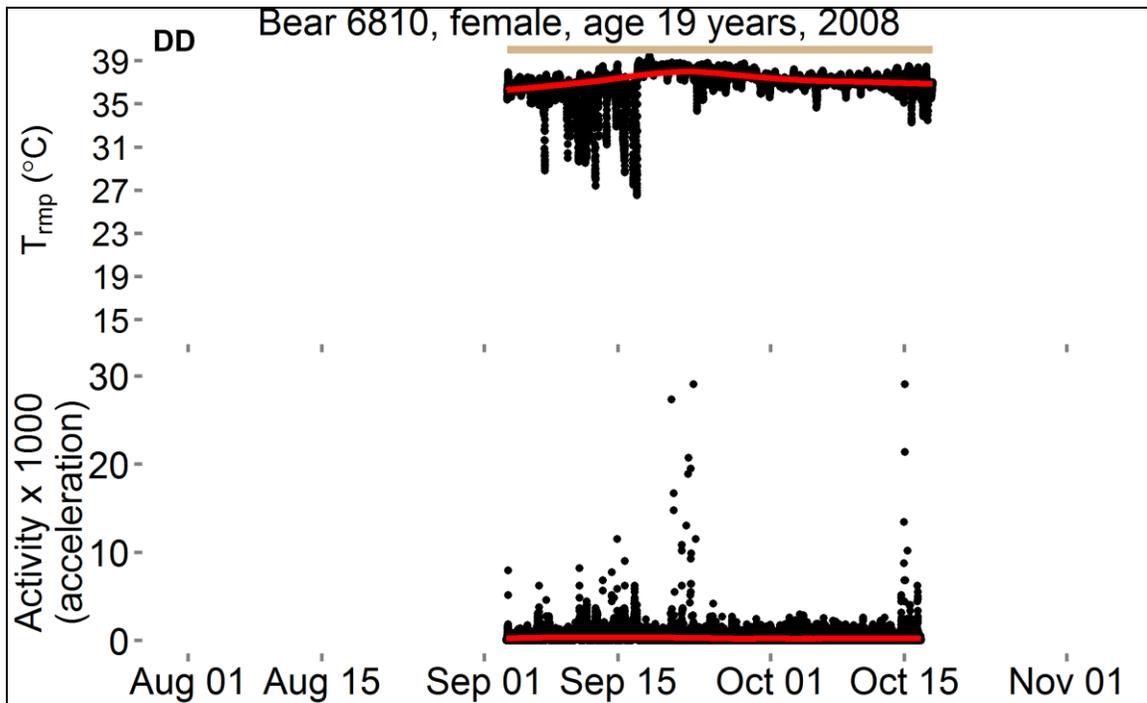


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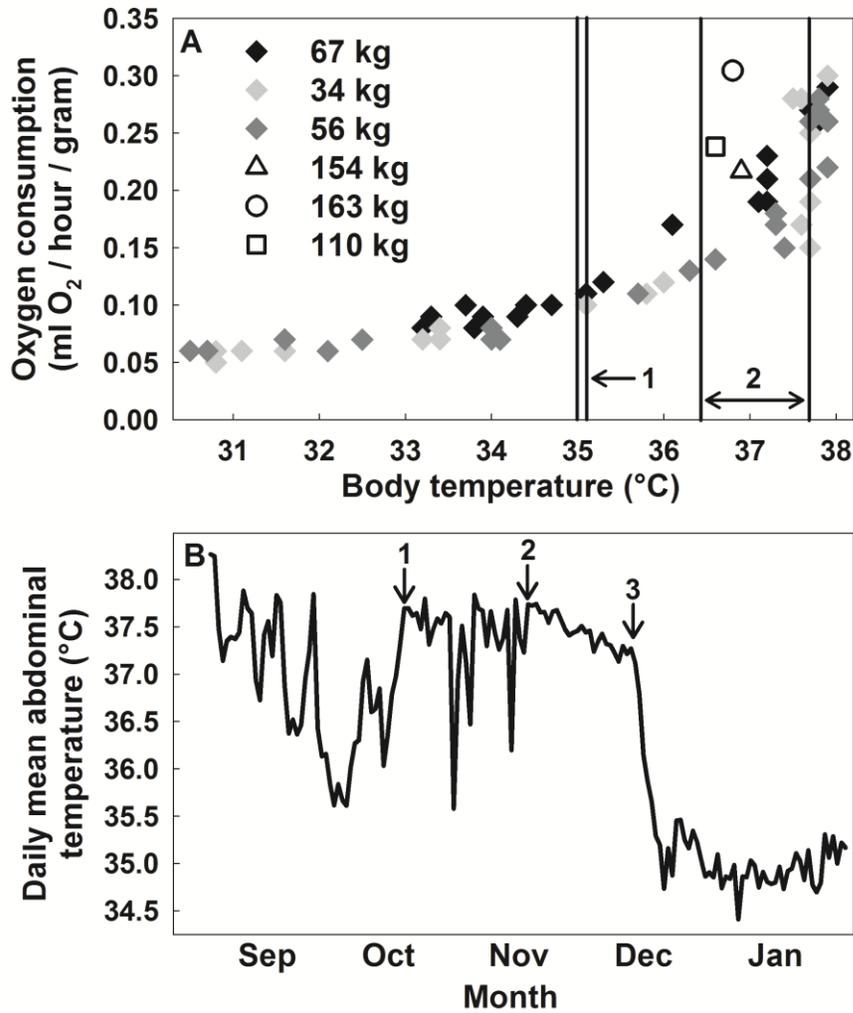
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351

**Fig. S1**

352 **Activity and body temperatures of polar bears in the Beaufort Sea, 2008–2010.** Each  
353 panel shows data from 1–3 variables for a single bear. Data censored  $\leq 5$  days after  
354 capture and  $\leq 1$  hour before recapture. Temperature is either “ $T_{abd}$ ” (hourly temperature  
355 of a logger implanted into the abdomen) or “ $T_{rmp}$ ” (temperature of a logger implanted  
356 beneath subcutaneous adipose tissue on the rump, recorded every 5–10 minutes), and red  
357 line is the smoothed trend after seasonal decomposition analysis. Activity is either  
358 “Activity (seconds)” (number of seconds of activity in previous half-hour) or “Activity  
359 (acceleration)” (acceleration score recorded every two minutes), and red line is moving  
360 average at center of 20-day window. For some bears, a horizontal bar spans the dates  
361 during which bear location was unknown (gray) or was on shore except for short swims  
362 (brown). The absence of a bar indicates the bear was offshore (swimming to or from sea  
363 ice, or traveling on sea ice surface). Data from bear 20741 were presented in Durner et al.  
364 (6).  
365

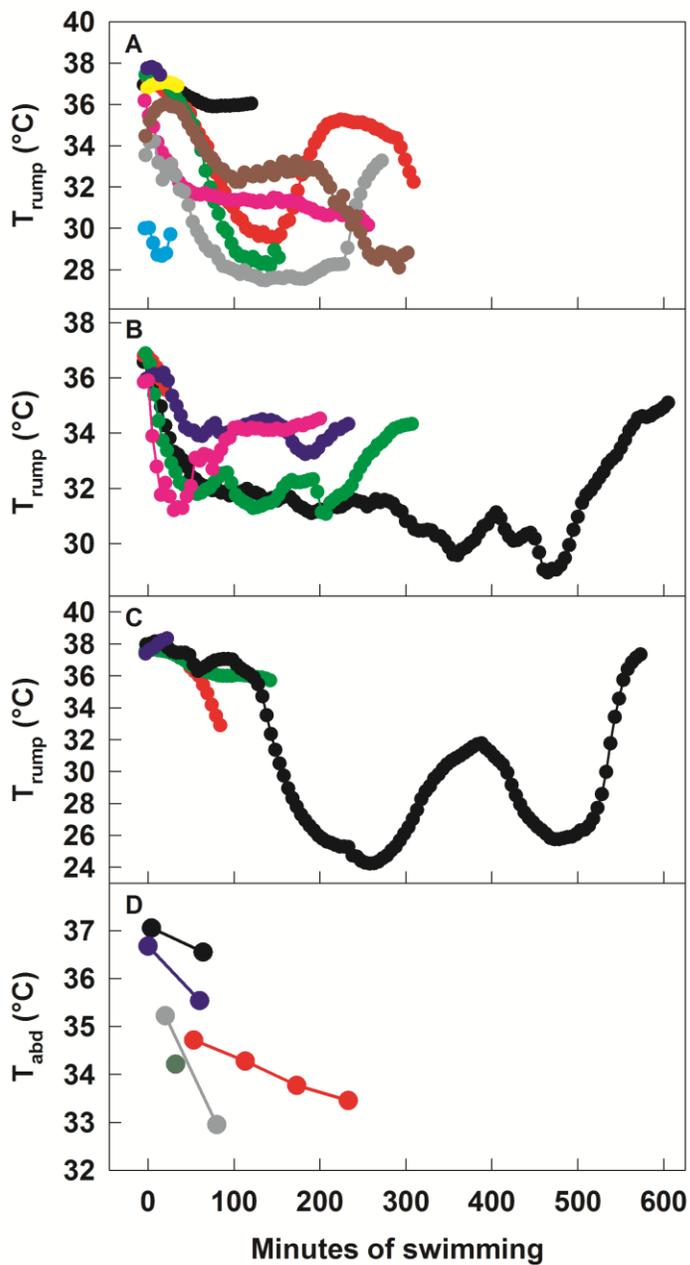


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368 **Fig. S2**

369 **Metabolic rates and body temperatures of bears.** (A) Data from previous studies  
 370 showing resting metabolic rates as measured by oxygen consumption of black bears  
 371 (solid diamonds; temperature logger inside peritoneum) and polar bears (open symbols;  
 372 temperature logger adjacent to peritoneum) of varying body mass (data adapted from  
 373 Hurst et al. (38) and Tøien et al. (27)). The spans on the x-axis indicate the range of  
 374 monthly temperature means from polar bears in our study for December–January (span 1;  
 375 bear 20529) and May–October (span 2; all other bears). (B) Abdominal temperatures of  
 376 pregnant polar bear 20529 in our study. At 1, an abrupt increase in her abdominal  
 377 temperature likely indicated blastocyst implantation, followed by entry into a maternity  
 378 den (at 2) and parturition (at 3).

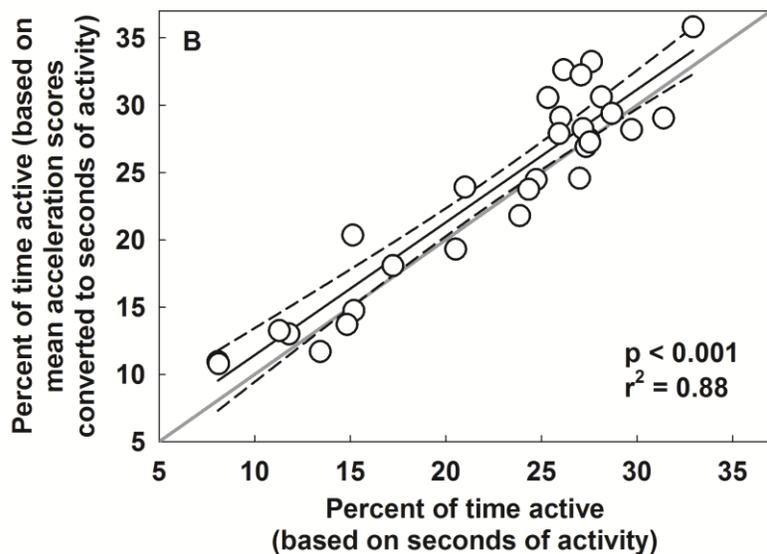
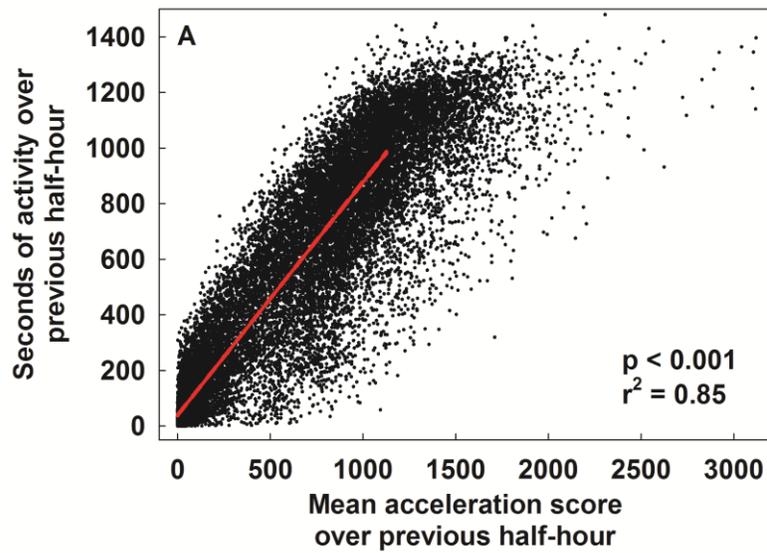
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382

383 **Fig. S3**

384 **Polar bear temperatures while swimming.** Data recorded by loggers implanted into the  
385 rump ( $T_{\text{rump}}$ ; A–C; measurements every five minutes) or abdomen ( $T_{\text{abd}}$ ; D;  
386 measurements hourly). Each panel represents a single bear and each color represents a  
387 single swim that was confirmed with a combination of location and activity data.  
388

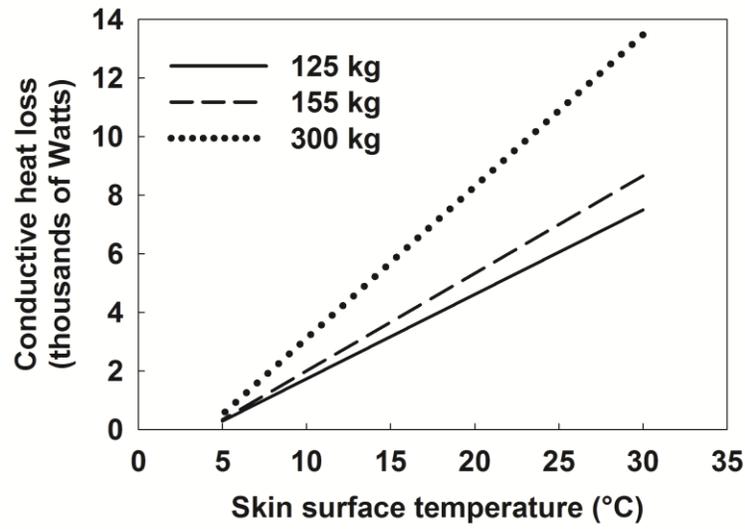


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390

391 **Fig. S4**

392 **Relationship between data collected simultaneously by two different collar-mounted**  
 393 **accelerometers on seven polar bears.** (A) Data are pooled across all bears. Statistics  
 394 describe the linear regression (solid red line; 95% confidence interval is narrow and  
 395 cannot be distinguished from the line itself) which extends from x values of 0 to 1128  
 396 (the cutoff identified by segmented regression). (B) The x-axis represents a monthly  
 397 mean derived from number of seconds of activity per half-hour, and the y-axis represents  
 398 a monthly mean derived from acceleration scores converted to number of seconds of  
 399 activity per half-hour. The relationship  $x=y$  is also shown (gray line). Statistics describe  
 400 the linear regression (solid black line; 95% confidence interval shown by dashed black  
 401 lines).

402



403  
404

405 **Fig.S5**

406 **Estimated heat loss of swimming polar bears.** Heat loss is influenced by bear body  
407 mass (different lines) and skin surface temperature.

408  
409

410 **Table S1.**  
 411 **Statistics of Welch t-tests comparing data collected from polar bears on shore and**  
 412 **on sea ice.** Where present, data from 2008 and 2009 were combined. The variable of time  
 413 spent active is derived from both seconds of activity per half-hour, and from acceleration  
 414 scores converted to seconds of activity per half-hour.  
 415

Variable	Month	Mean $\pm$ SE		<i>p</i>	t	df
		Shore	Ice			
Time spent active	July	11.8	24.6 $\pm$ 0.9	NA	NA	NA
	August	21.4 $\pm$ 5.3	20.0 $\pm$ 1.9	0.80	0.26	7.50
	September	17.8 $\pm$ 2.1	14.6 $\pm$ 1.1	0.21	1.33	13.54
	October	19.2 $\pm$ 2.7	19.3 $\pm$ 1.8	0.99	-0.01	12.71
Movement rate	July	0.12	0.31 $\pm$ 0.02	NA	NA	NA
	August	0.16 $\pm$ 0.02	0.32 $\pm$ 0.01	<0.01	7.55	9.85
	September	0.11 $\pm$ 0.02	0.33 $\pm$ 0.03	<0.01	-6.34	19.14
	October	0.14 $\pm$ 0.04	0.40 $\pm$ 0.05	<0.01	-4.35	17.99
Abdominal temperature	August	37.0 $\pm$ 0.2	36.8 $\pm$ 0.1	0.42	-0.88	4.50
	September	36.8 $\pm$ 0.2	36.6 $\pm$ 0.1	0.48	-0.74	6.24
	October	37.1 $\pm$ 0.2	36.8 $\pm$ 0.1	0.21	-1.44	5.13

416  
 417

418 **Table S2.**  
 419 **Statistics of Welch t-tests comparing data collected in 2008 and 2009, from polar**  
 420 **bears on shore.** Polar bears on sea ice were only sampled in 2009. The variable of time  
 421 spent active is derived from both seconds of activity per half-hour, and from acceleration  
 422 scores converted to seconds of activity per half-hour.  
 423

Variable	Month	Mean $\pm$ SE		<i>p</i>	<i>t</i>	df
		2008	2009			
Time spent active	July	NA	11.8	NA	NA	NA
	August	28.4 $\pm$ 7.5	12.2 $\pm$ 2.7	0.12	2.03	3.73
	September	18.6 $\pm$ 2.5	15.7 $\pm$ 4.5	0.60	0.58	3.33
	October	18.7 $\pm$ 3.1	22.7	NA	NA	NA
Movement rate	July	NA	0.12	NA	NA	NA
	August	0.16 $\pm$ 0.03	0.16 $\pm$ 0.02	NA	NA	NA
	September	0.10 $\pm$ 0.02	0.16 $\pm$ 0.03	0.26	-1.50	2.17
	October	0.14 $\pm$ 0.5	0.12	NA	NA	NA

424

425 **Table S3.**  
 426 **Coefficients from ARIMA models using environmental variables to predict daily**  
 427 **mean female polar bear activity.** Activity was measured as seconds of activity in the  
 428 previous half-hour, or as mean acceleration score over the previous half-hour. Sample  
 429 size indicates the number of days upon which both variables were measured. Coefficients  
 430 which overlapped with zero indicated no significant relationship. Variables include  
 431 “Shelf” (binary: whether mean depth of water for bears on sea ice was  $\leq 300$  m deep),  
 432 “Whale” (daily proportion of hourly locations  $< 500$  from whaling site), and “Air temp”  
 433 (daily mean of air temperature at nearby weather station).  
 434

Response	Predictor	Bear ID	Coefficient estimate (95% CI)	<i>n</i>
Seconds of activity per half- hour	Shelf	20157	1.01 (-3.16 – 5.17)	106
		20414	0.10 (-3.58 – 3.77)	96
		20446	0.05 (-1.88 – 1.97)	95
		20474	0.94 (-2.74 – 4.63)	36
		20479	-8.02 (-14.72 – -1.32)*	176
		20520	0.96 (-2.59 – 4.52)	133
		20534	2.62 (0.64 – 4.60)*	161
		20735	-2.08 (-3.91 – -0.24)*	72
		20764	-0.46 (-2.16 – 1.24)	371
		20901	0.90 (-2.58 – 4.38)	141
		21024	1.33 (-0.85 – 3.50)	324
		21032	2.19 (-2.12 – 6.50)	139
		21035	3.88 (2.30 – 5.46)*	104
		21045	-1.79 (-3.33 – -0.25)*	315
32777	0.21 (-1.81 – 2.22)	156		
<i>Combined weighted mean coefficient (95% CI)</i>			<i>0.43 (-0.16 – 1.02)</i>	
Acceleration score	Shelf	20414	-0.36 (-1.18 – 0.46)	66
		20446	-0.05 (-0.29 – 0.19)	61
		20735	-0.17 (-0.35 – 0.02)	56
		20741	-0.29 (-0.91 – 0.33)	56
		20764	0.03 (-0.09 – 0.14)	41
		21024	0.30 (-0.03 – 0.64)	92
		21045	-0.20 (-0.42 – 0.02)	181
		32777	0.17 (-0.17 – 0.51)	92
<i>Combined weighted mean coefficient (95% CI)</i>			<i>-0.03 (-0.11 – 0.05)</i>	
Acceleration score	Whale	20586	0.00 (-0.26 – 0.27)	50
		6810	-0.50 (-0.92 – -0.07)*	42
		20975†	-3.33 (-7.88 – 1.22)	27
		32282	2.69 (0.98 – 4.41)*	39
		32606	0.25 (-0.45 – 0.94)	61
		20966	0.78 (0.01 – 1.55)*	64

<i>Combined weighted mean coefficient (95% CI)</i>			<i>-0.01 (-0.20 – 0.22)</i>	
Seconds of activity per half-hour	Whale	20735	10.30 (-8.17 – 28.77)	72
<i>Combined weighted mean coefficient (95% CI)</i>			<i>0.00 (-0.01 – 0.01)</i>	
Acceleration score	Air temp.	20414	0.14 (-0.01 – 0.30)	30
		20446	0.01 (-0.11 – 0.12)	31
		20586	0.00 (-0.02 – 0.02)	50
		20735	0.05 (-0.02 – 0.11)	33
		20966	0.00 (-0.03 – 0.02)	64
		20975†	0.05 (0.02 – 0.08)*	27
		32255	-0.08 (-0.12 – -0.04)*	36
		32282	0.02 (-0.01 – 0.04)	39
		6810	0.11 (-0.19 – -0.02)*	42
<i>Combined weighted mean coefficient (95% CI)</i>			<i>0.00 (-0.01 – 0.01)</i>	
Seconds of activity per half-hour	Air temp.	20414	0.53 (-0.17 – 1.23)	30
		20446	-0.21 (-0.55 – 0.13)	41
		20735	-0.17 (-0.40 – 0.05)	70
<i>Combined weighted mean coefficient (95% CI)</i>			<i>-0.14 (-0.22 – 0.04)</i>	

435 \* Does not overlap with zero

436 †Male

437

438

439 **Table S4.**  
 440 **Hourly locations recorded from GPS telemetry transmitters on female polar bears**  
 441 **on shore between August 1<sup>st</sup> and November 1<sup>st</sup>.** Data from 2008 and 2009 were pooled.  
 442 Expected total number of locations is the number of hours between capture and recapture  
 443 of bears. Actual number of locations is fewer than expected because weather and  
 444 equipment failure occasionally prevented transmitter function. For each bear, a total of  
 445 the number of locations that were within 2000 m and 500 m of three sites where bowhead  
 446 whale carcasses are deposited after human harvest is also presented.  
 447

BearID	Year	Number of hourly locations			
		Expected total	Actual total (% of expected)	Within 2000 m (% of actual)	Within 500 m (% of actual)
32282	2008	1103	1099 (99)	431 (39)	41 (4)
20735	2009	1677	1666 (99)	885 (53)	59 (4)
32255	2008	1030	1017 (99)	0 (0)	0 (0)
20966	2008	1700	1675 (98)	898 (54)	107 (6)
20586	2008	1386	1363 (98)	1084 (80)	931 (68)
20446	2009	945	926 (98)	14 (2)	1 (0.1)
6810	2008	1193	1144 (96)	701 (61)	649 (57)
32608	2008	1193	1041 (87)	537 (52)	50 (5)
20982	2008	1517	1104 (73)	242 (22)	28 (3)
32606	2008	402	279 (69)	155 (56)	23 (8)
20965	2008	1611	1031 (64)	159 (15)	20 (2)
20974	2008	1732	899 (52)	0 (0)	0 (0)
20414	2009	1909	930 (49)	0 (0)	0 (0)
20492	2008	1823	818 (45)	386 (47)	78 (10)
20975*	2008	1659	685 (41)	307 (45)	25 (4)

448 \*Male  
 449

450 **Table S5.**  
 451 **Correlations between hourly body temperatures and hourly activity of polar bears**  
 452 **in the Beaufort Sea, estimated with dynamic linear models.** Both body temperature  
 453 variables were filtered to include only data < 4 SD from the smoothed mean. For rump  
 454 temperature, data also filtered to only include measurements recorded when bears were  
 455 active for > 2% of the previous half-hour.  
 456

Variables		Bear ID	Correlation (95% CI)	<i>n</i>
Body temperature	Activity	20414	0.24 (0.16 – 0.32)	703
		20520	0.17 (0.13 – 0.21)	2261
		20764	0.38 (0.35 – 0.41)	3595
		21045	0.46 (0.43 – 0.49)	3375
		32777	0.34 (0.31 – 0.37)	3623
Abdominal temperature	Seconds of activity per half-hour	20414	0.21 (0.17 – 0.25)	703
		20764	0.21 (0.15 – 0.27)	977
		21045	0.50 (0.46 – 0.54)	2181
		32777	0.25 (0.21 – 0.29)	2181
		21024	0.26 (0.22 – 0.30)	2995
Rump temperature	Acceleration score	20586	0.26 (0.20 – 0.32)	754
		32255	0.32 (0.22 – 0.42)	473
		32282	0.40 (0.34 – 0.47)	681
		6810	0.11 (0.03 – 0.19)	642
		20741	0.03 (-0.05 – 0.11)	966

457

458 **Table S6.**  
 459 **Body mass of polar bears.** Measurements taken on the date of implantation of  
 460 temperature loggers and at the date of logger retrieval (dates in 2009, except as noted).  
 461

Bear	Implant		Retrieval	
	Date	Body mass (kg)	Date	Body mass (kg)
Abdominal loggers				
20132	April 29	160	April 22*	148
20414	April 24	208	October 10	243
20520	May 09	179	April 17*	229
20529	August 11	350	April 05*	172
20562	August 04	238	October 07	211
20764	May 01	207	October 03	210
20947	August 10	434	October 11	465
21045	May 19	263	October 12	299
21150	August 10	123	October 18	189
32777	May 09	196	October 19	299
Rump loggers				
20586	August 22†	270	October 19†	323
20741	August 23†	226	October 26†	177
20898	August 16†	168	NA‡	NA‡
21024	April 25	179	October 07	225
32255	August 26†	218	October 08†	187
32282	August 25†	273	October 10†	291
6810	August 28†	234	October 17†	288

462 \*2010  
 463 †2008  
 464 ‡Logger retrieved from harvested bear  
 465

466 **Table S7.**  
 467 **Sample size (number of polar bears) for three variables shown in Fig. 2.**  
 468

Month (habitat)	Percent time active	Movement rate (m/s)	Proportion of movement rates > 0.33 m/s recorded when bears were inactive
April (ice)	2	2	2
May (ice)	15	15	15
June (ice)	15	15	15
July (ice)	14	15	13
July (shore)	1	1	1
August (ice)	10	9	9
August (shore)	7	5	4
September (ice)	12	13	8
September (shore)	10	11	7
October (ice)	10	11	9
October (shore)	8	9	3
November (ice)	3	3	3
December (ice)	3	3	3
January (ice)	3	3	3
February (ice)	3	3	3
March (ice)	3	3	3

469