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1	Post-fire recovery of torpor and activity patterns of a small mammal
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16	
17	Abstract
18	To cope with the post-fire challenges of decreased availability of food and shelter, brown
19	antechinus (Antechinus stuartii), a small marsupial mammal, increase the use of energy
20	conserving torpor and reduce activity. However, it is not known how long it takes for animals
21	to resume pre-fire torpor and activity patterns during the recovery of burnt habitat. Therefore,
22	we tested the hypothesis that antechinus will adjust torpor use and activity after a fire
23	depending on vegetation recovery. We simultaneously quantified torpor and activity patterns
24	for female antechinus from three adjacent areas: (i) the area of a management burn one year
25	post-fire, (ii) an area that was burned two years prior and (iii) a control area. In comparison to

shortly after the management burn, antechinus in all three groups displayed less frequent and less pronounced torpor while being more active. We provide the first evidence that only one year post-fire antechinus resume pre-fire torpor and activity patterns, likely in response to the return of herbaceous ground cover and foraging opportunities.

30

31 Keywords: antechinus, behaviour, heterothermy, marsupial, physiology, predation32

33 1. Introduction

As large and destructive wildfires are increasing worldwide [1], management burns are
employed in many forested areas with the aim of reducing fire fuel loads. However, the time
between fires and their intensity can alter assemblages of plant and animal species [2,3].
Importantly, while leaf litter and some grasses/herbs often reappear quickly after a fire,
others, such as woody vegetation and tree hollows, can take up to 30-40 years to re-establish
[4].

In the long-term fires can help to increase habitat complexity, which is beneficial for 40 many small terrestrial mammals [2]. However, the initial loss of ground cover after a fire can 41 42 be detrimental because it leads to reduced foraging opportunities due to increased predation pressure [5]. This is exacerbated for insectivorous mammals, as the abundance of ground-43 44 dwelling arthropods often decreases after fire [6]. Hence, in the short to medium-term after fire, some terrestrial mammals have been found to substantially curtail activity and enhance 45 the use of torpor - a pronounced reduction in energy expenditure via a controlled fall of 46 metabolic rate and body temperature (T_b) [6-9]. 47

Given the slow recovery of some habitat features it remains unclear over what time period such behavioural and physiological adjustments have to be employed for individuals and populations to survive. While an increase in torpor use is beneficial during detrimental

conditions, it may reduce fitness by curtailing territorial defence and reproduction. Therefore,
as conditions improve animals should employ less torpor. We tested the hypothesis that
following the recovery of an herbaceous understory after a management burn, torpor use and
activity in brown antechinus (*Antechinus stuartii*), a small heterothermic terrestrial marsupial
[10], will return to pre-fire levels in response to more profitable and safer foraging
opportunities.

57

58 2. Material and methods

59 Our work was undertaken during May-June 2015, one year after a management burn at Guy Fawkes River National Park, Australia (30°04'58.6"S, 152°20'0.9"E). We compared data 60 collected during the current study to those from our previous study in 2014 [9], with a total of 61 62 six groups of female antechinus (defined in table 1). All groups were measured in areas that were in close proximity, ensuring that the general habitat and climate of the sites was the 63 same. The areas that burnt in 2013 and 2014 are burnt regularly to reduce fuel levels and the 64 intensity of wildfires, whereas the control area last burned in 1994 due to a wildfire and no 65 management burns have been implemented since [11]. Ambient temperatures (T_a) were 66 recorded with data loggers ($\pm 0.5^{\circ}$ C, iButton thermochron DS1921G, Maxim Integrated 67 Products, Inc., Sunnyvale, California, USA). We visually determined the percentage of 68 ground cover in 12 randomly selected 1m² quadrats in each of the three study areas (36 69 70 quadrats total).

To measure T_b patterns including torpor and also to infer activity we implanted
temperature-sensitive radio-transmitters (2g, Sirtrack, Havelock North, New Zealand) into the
intraperitoneal cavity of each animal (for surgical details see [12]). A receiver/data logger
[13] was placed near each nest to record the presence/absence of animals and their T_b.
Throughout the study each individual used between 1 and 8 nest sites (mean = 3.7 nests) and

loggers were repositioned whenever antechinus moved to a new nest. Activity periods within
24 h (sunrise to sunrise) were calculated as the time animals spent away from the nest
site/logger for >30min. Days when an individual moved nests were excluded as the time the
animal arrived at the new nest was unknown. We calculated the torpor threshold of 31.5°C
using equation 4 [14] as previously [9]; torpor bouts were calculated from the time T_b fell
below and rose above this threshold for > 30 min.

82 We used R [15] to perform statistical analyses. As the duration of activity and torpor are significantly correlated with T_a in antechinus [10], only data recorded when T_a was within 83 84 daily maxima (20.0°C) and minima (2.0°C) of our previous study were included [9]. We employed linear mixed-effects models [16] to test for differences among the six groups for 85 the measured variables: torpor frequency, torpor bout duration, daily minimum T_b, time spent 86 active [17]. We used body mass and daily minimum T_a as covariates, individuals were 87 included as a random effect to account for repeated measures and we accounted for temporal 88 autocorrelation [17]. We used a residual plot to test for homoscedasticity and a normal Q-Q 89 plot to test for normal distribution. We employed a post-hoc Tukey test [18] to determine 90 which groups differed from each other. All proportions were arcsine transformed before 91 analysis. We assumed significance at p < 0.05. 92

93

94 **3. Results**

While ground cover had somewhat recovered a year after the burn in 2014 (figure 1), it was patchy in those areas where the one year post-fire 2015 (range = 10-100%, mean = 50%, n =12) and two years post-fire 2015 (range = 5-100%, mean = 39%, n = 12) groups were measured. In contrast, the area where the control 2015 group was measured consisted of thick and even ground cover unchanged from 2014 (range = 60-100%, mean = 87%, n = 12).

In all 2015 groups torpor was employed approximately half as often as by the postfire 2014 group, and torpor frequency was therefore statistically identical to the pre-fire 2014 and control 2014 groups (figure 2*a*; table 2). Furthermore, in all 2015 groups mean torpor bout durations and the longest torpor bouts recorded per individual were significantly shorter (~50%) compared to the post-fire 2014 group (figure 2*b*; table 2). The mean daily minimum T_b displayed by groups in the 2015 study was ~3°C higher, and the lowest T_b recorded >3.4°C greater, than that of the post-fire 2014 group (figure 2*c*; table 2).

107 Antechinus from all three groups measured in 2015 spent an additional three hours 108 active each day in comparison to individuals from the post-fire 2014 group (figure 2*d*; table 109 2). Further, while daytime activity was almost absent in the post-fire 2014 group (4.4% of 110 total daily activity), it encompassed ~20% of the total daily activity of antechinus in 2015 and 111 of those in the pre-fire 2014 and control 2014 groups (figure 2*d*).

112

113 4. Discussion

Our study provides the first evidence that within one year after a management burn a population of brown antechinus had resumed pre-fire torpor and activity patterns. Torpor bouts were shorter and shallower in individuals in the post-fire recovered landscape in comparison to shortly after the fire when ground cover was largely absent. Further, this decrease in torpor use one year after the fire was accompanied by an increase in total and especially daytime activity over levels measured just after the burn.

The late autumn management burns employed for our study were of low-intensity, but inflicted some mortality amongst antechinus nevertheless [9]. As the fires primarily consumed dead vegetation and dry leaf litter leaving the mid and upper storeys largely intact, a relatively quick recovery of some vegetative ground cover ensued. In particular *Lomandra* spp. (native Australian perennial herbs) had re-sprouted vigorously in the areas that had

burned both one and two years prior (figure 1). However, *Lomandra* is encouraged by fire,
and other types of ground cover such as small shrubs and decaying timber were still largely
absent. Yet, brown antechinus persisted and during early winter resumed daily torpor and
activity patterns similar to those recorded at the same time of year before the fire and in
unburnt habitat [9].

Firstly, it therefore appears that even sparse vegetation of limited diversity is 130 131 sufficient for antechinus to manage their daily energy needs without extensive torpor use. This vegetative cover likely aids in protecting antechinus from predation while foraging. 132 133 Secondly, as the abundance of terrestrial arthropods often increases in the year following a fire [6], for antechinus the prey base might have been elevated. Thirdly, it is likely that the 134 insulating properties of a tussock ground cover alters the microclimate of foraging areas. For 135 example, the retention of a boundary layer of warm air near the ground during the night 136 would reduce thermoregulatory costs while foraging, further decreasing the need to save 137 energy via torpor. 138

Our results highlight the importance of even sparse ground cover for small terrestrial 139 mammals while foraging [19]. The general association of torpor with low T_a and the 140 reduction of torpor use as habitat conditions improve suggests that antechinus mostly employ 141 torpor in response to detrimental conditions, which is similar to other marsupials such as 142 sugar gliders (*Petaurus breviceps*) [9,10,13]. Importantly, antechinus are somewhat unusual 143 144 in being semelparous, reproducing only once a year towards the end of winter [20]. It is therefore likely vital for them during autumn and early winter to invest time in protecting 145 resources, such as food and nests, to ensure reproductive success, and torpor decreases the 146 time available for these activities. 147

148 Nevertheless, it is evidently possible that fires do not pose a significant challenge for
149 some mammals, especially those that can avoid fires like volant bats [21]. Indeed, the only

150	other study we are aware of focussing on how individuals recover physiologically and
151	behaviourally in the years after a fire was conducted on bats [21]. Therefore, further studies
152	on different mammals are needed to understand how other species on other continents
153	respond physiologically and behaviourally to fires, including severe wildfires, in the short- to
154	long-term, particularly on longer-lived and homeothermic species that do not have the ability
155	to employ torpor for energy conservation.
156	
157	Ethics. Permits for this study were issued by the University of New England Animal Ethics
158	Committee (AEC13-088) and the New South Wales National Parks and Wildlife Service
159	(SL100791).
160	
161	Data accessibility. Data can be accessed from Dryad Digital Repository
162	(doi:10.5061/dryad.m76ks).
163	
164	Authors' contributions. C.S. and F.G. designed the study. C.S. and T.H. analysed the data.
165	C.S. drafted the manuscript. All authors collected the data and commented on the manuscript.
166	All authors agree to be held accountable for the content.
167	
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169	
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225

Figure 1. Photos of the study site depicting ground cover in the same area shortly after the management burn in 2014 (*a*) and one year later in 2015 when *Lomandra* had recovered (*b*).



229

Figure 2. Comparison of variables for female *Antechinus stuartii* from 2014 (white bars) and 2015 (grey bars). (*a*) torpor frequency (number of days with torpor/days measured), (*b*) mean torpor bout duration (min), (*c*) mean daily minimum T_b (°C) and (*d*) time spent active (min). For activity (*d*) the solid portion of the bar represents daytime (sunrise-sunset) activity and the hatched portion nocturnal (sunset-sunrise) activity. Means are shown ± 1 standard deviation, which for (*d*) represents whole day (24h, sunrise-sunrise) activity. Only the postfire 2014 group differed significantly, denoted by the asterisk.

Table 1. Descriptions of each of the groups measured, including number of individuals and

238	animal	days	measured.

group name	group description	individuals	animal days
pre-fire 2014	individuals measured in 2014 before the burn	4	25
post-fire 2014	individuals measured in 2014 after the burn	4	56
control 2014	individuals measured in 2014 in an unburnt area	2	17
one year post-fire 2015	individuals measured in 2015 one year after the management burn in 2014	7	136
two years post-fire 2015	individuals measured in 2015 in an area that was burnt two years prior in 2013	4	89
control 2015	individuals measured in 2015 in an unburnt area	6	135

Table 2. Torpor variables, body temperatures (T_b) and time spent active (whole day) of female *Antechinus stuartii* and ambient temperatures

241	(T _a). Means are shown ± 1	standard deviation and	l significant differen	ces between groups	are identified by dif	ferent letters.
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	pre-fire 2014	post-fire 2014	control 2014	one year post-fire 2015	two years post-fire 2015	control 2015
torpor frequency (proportion of days)	$0.52\pm0.27^{\rm a}$	0.92 ± 0.07^{b}	$0.43\pm0.35^{\mathtt{a}}$	$0.57\pm0.22^{\mathtt{a}}$	$0.62\pm0.16^{\rm a}$	$0.50\pm0.28^{\rm a}$
torpor bout duration (min)	$120.8\pm80.8^{\text{a}}$	274.2 ± 79.2^{b}	$137.5\pm102.5^{\text{a}}$	142.6 ± 81.4^{a}	$126.7\pm33.1^{\text{a}}$	$121.9\pm69.1^{\mathrm{a}}$
longest torpor bout recorded (min)	330	650	310	380	340	340
minimum T _b (°C)	$31.3\pm2.2^{\rm a}$	$27.4 \pm 1.9^{\text{b}}$	$31.2\pm2.8^{\rm a}$	$30.1 \pm 1.9^{\mathrm{a}}$	$29.5\pm1.4^{\rm a}$	$31.1\pm2.3^{\rm a}$
lowest T _b recorded (°C)	24.3	19.8	27.6	24.0	23.2	25.1
time spent active (min)	$693.2\pm62.7^{\mathtt{a}}$	$408.2\pm137.6^{\text{b}}$	$617.1\pm107.0^{\text{a}}$	$599.6\pm106.6^{\mathtt{a}}$	$661.2\pm68.1^{\text{a}}$	$629.3\pm75.7^{\rm a}$
daily T _a (°C)	$8.3\pm2.8^{\rm a}$	11.1 ± 1.2^{b}	$10.9\pm1.1^{\text{b}}$	$9.6\pm1.1^{\text{b}}$	9.9 ± 1.0^{b}	$9.9\pm1.3^{\text{b}}$
maximum T _a (°C)	$13.5\pm2.6^{\rm a}$	16.5 ± 1.7^{b}	$16.3\pm2.0^{\text{b}}$	15.8 ± 0.8^{b}	$15.4\pm1.2^{\text{b}}$	$15.7\pm1.3^{\rm b}$
absolute highest maximum T _a (°C)	17.5	20.0	19.5	18	18.5	18
minimum T _a (°C)	$4.8\pm2.8^{\text{a}}$	$7.4 \pm 1.4^{\text{b}}$	7.0 ± 1.4^{b}	6.8 ± 1.5^{b}	7.1 ± 1.4^{b}	7.1 ± 1.7^{b}
absolute lowest minimum T _a (°C)	2.0	5.5	5.0	5.5	5.5	5.0