American Journal of Primatology, in press, 2019.

Individual, social and environmental factors affecting salivary and fecal cortisol levels in captive pied tamarins (*Saguinus bicolor*)

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Dedication: In memory of Tine Griede

Running head: Fecal and salivary cortisol in pied tamarins

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1 Abstract

2 Pied tamarins (Saguinus bicolor) are endangered New World primates, and in 3 captivity appear to be very susceptible to stress. We measured cortisol in 214 saliva 4 samples from 36 tamarins and in 227 fecal samples from 27 tamarins, and investigated 5 the effects of age, sex, pregnancy, rearing history, social status, weight, group 6 composition and enclosure type using generalized linear mixed models. There was no 7 effect of age on either fecal or salivary cortisol levels. Female pied tamarins in late 8 pregnancy had higher fecal cortisol levels than those in early pregnancy, or non-9 pregnant females, but there was no effect of pregnancy on salivary cortisol. Females 10 had higher salivary cortisol levels than males, but there was no effect of rearing 11 history. However, for fecal cortisol, there was an interaction between sex and rearing 12 history. Hand-reared tamarins overall had higher fecal cortisol levels, but while male 13 parent-reared tamarins had higher levels than females who were parent-reared, the 14 reverse was true for hand-reared individuals. There was a trend towards lower fecal 15 cortisol levels in subordinate individuals, but no effect of status on salivary cortisol. Fecal but not salivary cortisol levels declined with increasing weight. We found little 16 17 effect of group composition on cortisol levels in either saliva or feces, suggesting that 18 as long as tamarins are housed socially, the nature of the group is of less importance. 19 However, animals in off-show enclosures had higher salivary and fecal cortisol levels 20 than individuals housed on-show. We suggest that large on-show enclosures with 21 permanent access to off-exhibit areas may compensate for the effects of visitor 22 disturbance, and a larger number of tamarins of the same species housed close

- 23 together may explain the higher cortisol levels found in tamarins living in off-show
- 24 accommodation, but further research is needed.
- 25
- 26 **Key words:** callitrichid, cortisol, *Saguinus bicolor*, stress, welfare

27 28	Rese	earch highlights
29	•	Hand-reared pied tamarins have higher fecal cortisol levels than parent-reared
30		tamarins.
31	•	Pied tamarins living off-show with more conspecific groups per building have
32		higher salivary and fecal cortisol levels than tamarins housed on-show in large
33		enclosures with access to off-exhibit areas.

34

35 Introduction

36

37 Animal welfare science in zoos is a long-established field of study (Powell & 38 Watters, 2017). The highest standards of welfare in zoo-housed animals are essential, 39 particularly given the growing role of captive breeding programs in conservation 40 strategies for threatened species, and as zoos increasingly seek to educate as well as to 41 entertain their visitors (EAZA, 2013; Kagan, Carter, & Allard 2015). Zoos are also 42 challenged regularly about the ethics of keeping animals in captivity (Gross, 2015; 43 Keulartz, 2015), and thus appropriate measures of welfare are needed. Behavioral 44 studies are often used for this purpose (Dawkins, 2004), but measurements of 45 glucocorticoid hormones such as cortisol are increasingly common as a means of 46 assessing wellbeing (Möstl & Palme, 2002; Keay, Singh, Gaunt, & Kaur, 2006; Clark et 47 al., 2012; Hart, 2012). 48 As well as being influenced by factors such as social and reproductive status 49 (Abbott et al., 2003; Bales, French, Hostetler, & Dietz, 2005; Bales, French, McWilliams, 50 Lake, & Dietz, 2006), cortisol is produced by the hypothalamus–pituitary–adrenal axis 51 (HPA) when an individual is exposed to a stressful situation (Möstl & Palme, 2002; 52 Beehner & Bergman, 2017). Animals in natural environments can usually respond 53 adaptively (either behaviorally and or physiologically) to stressors and ameliorate the 54 stress (Morgan & Tromborg, 2007). Acute stress may therefore not have a negative 55 effect on fitness (Beehner & Bergman, 2017). However, if it is not possible to respond 56 in such a way as to reduce the effect of a stressor, as is often the case in animals living in captivity (Mason, 2010), stress can become chronic and the individual may develop 57

58	problems such as stereotypic behaviors (e.g. Mason, Clubb, Latham, & Vickery, 2007),
59	poor health (e.g. Munson et al., 2005), excessive weight loss (e.g. Tamashiro, Nguyen,
60	& Sakai, 2005), reduced reproductive success (e.g. Clubb, Rowcliffe, Lee, Mar, Moss, &
61	Mason, 2008), suppressed immunity (Martin, 2009), and impaired cognitive function
62	(Teixeira et al., 2015). A growing number of studies in a variety of mammal species
63	living in zoos has pointed to many factors that can affect cortisol levels, including
64	rearing history, social situation, enclosure size and type, noise or other disturbance
65	(e.g. from zoo visitors), access to outdoor areas, and season (Carlstead, Brown, &
66	Seidensticker, 1993; McCallister, 2005; Carlstead & Brown, 2005; Powell, Carlstead,
67	Tarou, Brown, & Monfort, 2006; Clark et al., 2012; Pirovino et al., 2011; Rajagopal,
68	Archunan, & Sekar, 2011; Cerda-Molina et al., 2012; Kaplan et al., 2012; Shepherdson,
69	Lewis, Carlstead, Bauman, & Perrin, 2013; Schumann, Guenther, Jewgenow, &
70	Trillmich, 2014; Sherwen et al., 2015; Pauling, Lankford, & Jackson, 2017).
71	Our study evaluates factors that might affect cortisol levels in the pied tamarin
72	(Saguinus bicolor), a callitrichid primate that is endemic to the Brazilian Amazonian
73	rain forest near the city of Manaus. Pied tamarins are classed by the IUCN as
74	endangered (Mittermeier, Boubli, Subirá, & Rylands, 2008), and their population
75	continues to decline because of severe habitat loss and fragmentation (Gordo, Calleia,
76	Vasconcelos, Leite, & Ferrari, 2013). Along with in-situ conservation measures, one of
77	the main goals of the conservation action plan for the species is to create a stable,
78	healthy captive population (ICMBio, 2011). However, this has been challenging as pied
79	tamarins appear to be particularly sensitive to the conditions of captivity (Wormell,
80	Brayshaw, Price, & Herron, 1996; Holm, Priston, Price, & Wormell, 2012; Armstrong &

81	Santymire, 2013). "Wasting syndrome", characterized by severe weight loss, diarrhea,
82	and alopecia, has been a particular problem in captive populations (laleggio & Baker,
83	1995; Wormell, 2000; Smithyman, 2012; Cabana, Maguire, Hsu, & Plowman, 2018).
84	Pied tamarins are also behaviorally different from other callitrichids in some respects,
85	e.g. in frequently giving birth during the day (Price, Payne & Wormell, 2016). Thus, to
86	adapt management and improve welfare, as much species-specific information as
87	possible is needed about the extent to which pied tamarins experience stress from
88	various sources, as assumptions based on information from other species may not be
89	valid.
90	The main aims of this research were therefore to determine which factors
91	influence cortisol levels in zoo-housed pied tamarins and to gain an insight into
92	potential sources of stress. We analyzed cortisol in feces and saliva, as samples can be
93	collected non-invasively (Queyras & Carosi, 2004; Heistermann, 2010), and may also
94	give different pictures of the hormonal status of an individual (Cook, 2012). Cortisol
95	reaches saliva in a matter of minutes after it is secreted and as such reflects an acute
96	response to an event. Fecal sampling of cortisol, however, represents a timescale of
97	hours or even days, and thus repeated fecal sampling may provide insight into
98	underlying, longer-term sources of variation in cortisol levels.
99	We had access to relatively large samples in a single institution, and so were able
100	to look at the effects of eight factors that might affect cortisol levels in this species,
101	including variables at the individual level (sex, age, weight, rearing history, pregnancy),
102	the social level (social status, group composition) and the environmental level
103	(enclosure type). We tested the following predictions:

104	1.	Cortisol levels in pied tamarins will increase with age: Previous research has
105		found variable effects of age on cortisol levels in primates, and in several
106		studies there is little or no relationship (e.g. Bergman, Beehner, Cheney,
107		Seyfarth, & Whitten, 2005; Bales et al., 2006; Pirovino et al., 2011; though see
108		Laudenslager, Jorgensen, & Fairbanks, 2012; Fourie, Jolly, Phillips-Conroy,
109		Brown, & Bernstein, 2015a). However, as older tamarins appear to be more
110		susceptible to wasting syndrome (Smithyman, 2012), we predicted that older
111		pied tamarins would have higher cortisol levels.
112	2.	Individuals with a higher body mass will have lower cortisol levels: Weight is
113		sometimes used as an indicator of stress (Schumann et al., 2014). We
114		therefore predicted a negative relationship between weight and cortisol levels.
115	3.	Cortisol levels will increase in female pied tamarins in late pregnancy:
116		Pregnancy has been found to affect cortisol levels in other callitrichid species,
117		with lower levels of cortisol in breeding females during early pregnancy, and
118		higher levels in late pregnancy (Bales et al., 2005; Ziegler, 2013).
119	4.	Female pied tamarins will have higher cortisol levels than males: Previous work
120		(Wark et al., 2016, Armstrong & Santymire, 2013) has found higher cortisol
121		levels in female callitrichids than in males.
122	5.	Hand-reared tamarins will have higher cortisol levels than parent-reared
123		individuals: Hand-rearing can have negative consequences for the behavior and
124		reproduction of primates in adulthood (Beck & Power, 1988; King & Mellen,
125		1994; Ryan, Thompson, Roth, & Gold, 2002), and hand-reared pied tamarins
126		often show higher aggression towards keepers than parent-reared animals

127		(Coe, 2014). They may therefore be more stressed by contact with people. As
128		hand-reared female pied tamarins never successfully rear their own infants
129		(Price et al., 2016), whereas a hand-reared male did become a competent
130		parent (pers. obs.), we also tested for an interaction between rearing and sex in
131		cortisol levels.
132	6.	Subordinate pied tamarins will not have higher cortisol levels than dominant
133		individuals: Tamarin groups vary considerably in composition and mating
134		system, but especially in captivity, are most often composed of a breeding pair
135		plus their offspring and sometimes other individuals (Anzenberger & Falk,
136		2012). The latter may remain in the group for considerable periods, but rarely
137		breed successfully (Price & McGrew, 1991; Savage, Giraldo, Soto & Snowdon,
138		1996; Saltzman, Liedl, Salper, Pick & Abbott, 2008; Henry, Hankerson, Siani,
139		French & Dietz, 2013). While in some primate species, subordinate status is
140		associated with higher cortisol levels (Abbott et al., 2003), several studies have
141		found that in the callitrichids, in which intragroup relationships are
142		predominantly affiliative rather than agonistic (Schaffner & Caine, 2000), either
143		dominant individuals have higher cortisol levels than subordinates, or there is
144		no effect of status (Ziegler, Scheffler, & Snowdon, 1995; Saltzman, Prudom,
145		Schultz-Darken, Wittwer, & Abbott, 2004; Bales et al., 2005, 2006).
146	7.	Tamarins living in mixed-species groups will have higher cortisol levels than
147		those living with conspecifics: Callitrichids live in close social groups with a
148		sophisticated cooperative rearing system (Tardif et al., 1986; Goldizen, 1987;
149		Price, 1992; Garber, 1997). Therefore, to avoid the potentially negative effects

150		of single housing, pied tamarins at Jersey Zoo that could not be housed with
151		conspecifics were usually mixed with individuals of other callitrichid species.
152		However, unlike some Saguinus species (Heymann & Buchanan-Smith, 2000),
153		pied tamarins are not usually sympatric with other callitrichids, and although
154		pied tamarins can be housed successfully with other species (pers. obs.), this is
155		not always the case (e.g. Gentry & Margulis, 2008).
156	8.	Tamarins housed in enclosures on show to the public will have higher cortisol
157		levels than tamarins living in off-show enclosures: Several studies have shown
158		that the presence of visitors increases the stress levels of both wild and zoo
159		animals (Hosey, 2000; Shepherdson, Carlstead, & Wielebnowski, 2004; Behie,
160		Pavelka & Chapman, 2010; Quadros, Goulart, Passos, Vecci, & Young, 2014;
161		Fourie et al., 2015b), and callitrichids in free-ranging environments that could
162		retreat further from the public had lower urinary cortisol levels than caged
163		conspecifics (McCallister, 2005).

164 Methods

165 Subjects and management

A total of 42 pied tamarins, all housed at Jersey Zoo in the Channel Islands, took part in the study (see Table 1); some individuals were included in studies of both salivary and fecal cortisol, while others contributed to only one data set. All tamarins involved in the present study were deemed healthy at the time the samples were collected, and were in stable social situations.

171	Tamarins all had permanent access to large indoor and outdoor areas,
172	predominantly in buildings housing 3–5 callitrichid groups (Wormell & Brayshaw,
173	2000). Indoor cages were of broadly similar size; minimum dimensions were
174	approximately 2.25 m high x 1.53 m wide x 2.45 m deep. All indoor areas received
175	natural light via skylights or windows. In addition, artificial lighting was provided via
176	strip lights and heat lamps from 0800 to 1800. In the winter months, supplementary
177	UV lighting was also put in place (López, Wormell, & Rodríguez, 2001). Outside
178	enclosures were 16–63 m ² in area and approximately 4 m high, and were planted with
179	extensive natural vegetation as well as being furnished with ropes, branches and
180	platforms. The design of the buildings meant that tamarins had no visual contact while
181	in their indoor areas, and no or very limited visual contact outside. Levels of auditory
182	and olfactory contact were similar in all buildings.
183	Three buildings were on show to the public, but only the outdoor areas were
184	accessible to visitors, and there were standoff barriers averaging 1 m from the front of
185	each enclosure, reducing the opportunity for visitors to touch the animals or the mesh
186	cage fronts. Three other buildings were off-show.
187	Pied tamarins were fed three times daily, at approximately 0800–0830 (primate
188	pellet mix), 1200–1300 (fruit, vegetables and a protein item such as egg), and 1530–
189	1700 (insect feed). Tamarin enclosures were cleaned in the morning, and excess food
190	removed in the late afternoon. Tamarins were trained to sit on scales within their
191	enclosures and were weighed at least weekly.

192

193 Sample collection

194 Salivary cortisol sampling and analysis

195	Saliva samples were collected between January and June 2007 by keeping staff
196	using the technique described by Cross, Pines, & Rogers (2004). Samples were
197	collected on a weekly basis. All samples included in the analysis were collected in the
198	morning, as cortisol levels decrease during the day (Cross & Rogers, 2004). Individuals
199	were encouraged to chew on 1–2 cotton buds for up to a minute at a time to obtain
200	the required volume of saliva (50 μ l); a single food incentive, honey, was used on the
201	cotton bud. The cotton buds were then centrifuged at 3200 rpm to extract the saliva
202	and frozen at –20° C until analysis by the Central Science Laboratory in York, UK, using
203	commercially available ELISA test kits and previously described methods (Cross et al.,
204	2004; Cross & Rogers, 2004; Gladwell & Pick, 2007).
205	

206 Fecal cortisol sampling and analysis

Fecal samples were collected between May and November 2008 at the first check in the morning (between 0800 and 0900), to control for circadian variation. Beetroot juice was used as a fecal marker and was given to tamarins via syringe in the evening prior to fecal deposition and collection the next morning. The animals were not disturbed by sample collection. Samples were frozen at -20°C within 60 min of collection.

213	Each sample was dried in a fan-assisted oven at 55°C for 7.5 h and was refrozen
214	until needed for further extraction at –20°C. Following thawing, each dry sample was
215	ground with a pestle and sifted through a fine wire mesh to remove seeds and fibrous
216	material (Wasser et al., 1993). A 3ml aliquot of 90% (v/v) methanol was added to a 0.1
217	g portion of the resulting powder and mixed vigorously for 3 h (Heidolphtitramax 100,
218	1350 rpm, 1.5 mm orbit). This was then centrifuged at 2000 rpm for 15 min following
219	an adapted version of the methods used by Wasser et al. (1993). The supernatant was
220	poured into a glass test tube and the ethanol evaporated using compressed oxygen
221	free nitrogen gas (N_2) administered using a Pierce Reacti-Therm Heating Module at
222	40°C. The residue was resuspended in 0.5 ml phosphate buffer saline buffer containing
223	0.1% (w/v) Bovine Serum Albumin (BSA) and microcentrifuged for 2 min at 6500 rpm
224	to remove any remaining solid particles. The resulting supernatant was stored at –20°
225	C until needed for measurement. Hormone values were expressed as $pg/50\mu l$ of fecal
226	extract.

227 Cortisol enzyme immunoassay

Immunological validation: A modified version of an enzyme immunoassay
described by Armstrong & Santymire (2013) was used to quantify levels of fGC. The
assay was immunologically validated for quantification of fGC levels in our population
of captive pied tamarins using a representative sample pool consisting of 50µL of
extract taken from all samples (Diamandis & Christopoulos, 1996). The antibody
(R4866, raised against a steroid bovine albumin in rabbit (Munro & Stabenfeldt, 1985))
was diluted to 1:12000 in coating buffer, and the cortisol horseradish peroxide was

235 diluted to 1:22000 in phosphate buffer solution containing 0.1% (w/v) BSA. Samples
236 were run in duplicate at 1: 100 dilution.

237 Cross-reactivity of the cortisol antibody was 100% with cortisol. Cross-reactivity 238 with similar steroids was 9.9% with prednisolone, 6.3% with prednisone, 5.0% with 239 cortisone, 0.7% with corticosterone and <0.3% with various other steroids (Munro & 240 Stabenfeldt, 1985). Assay specificity was demonstrated twice by parallel displacement 241 curves of serial dilutions of cortisol standard and the pied tamarin pool over the 10-242 90% binding range (ANCOVA; F_{3, 48} = 1.074, n.s.; F_{3, 41} = 1.053, n.s.). Recovery of the 243 standards (halving dilutions in the range from 500 to 7.8 pg) added to a 1:100 dilution of a mixed fecal pool was 96.16 \pm 14.91% inferring good accuracy (r = 0.996, F_{1.5} = 244 245 595.63, P < 0.0001). Intra-assay coefficients of variation for low, medium and high 246 concentration quality controls were 1.95, 4.79 and 4.93%, respectively. Interassay 247 coefficients of variation for low and high concentration quality controls were 10.03% 248 (n=5 plates) and 18.55% (n=7 plates). The sensitivity of the assay was approximately 249 1.95 pg/ml.

Biological validation: To determine whether our assay detected biologically meaningful changes, we assessed fGC levels in the morning and afternoon under control conditions to test for circadian variation this hormone. Samples were collected in the morning before 1100 (n = 83) and in the afternoon after 1400 (n = 97) from a mix of adult males and females over a 5-month period. Log transformed cortisol levels were compared between samples deposited in the morning versus the afternoon using an independent t test. Levels of fecal cortisol were significantly raised in the afternoon

compared to the morning in accordance with the typical circadian variation in levels of
excreted cortisol (t = 5.128, d.f. = 179, P < 0.001; mean ± S.E.M in the morning: 564.76
± 116.08 ng/ml and the afternoon: 1027.37 ± 117.03 ng/ml; Sousa & Ziegler, 1998).

261 Data analysis

262 General information about the samples, including dates, times, individuals, and 263 any events such as illness, social tension or catch-ups that might affect stress levels, 264 was recorded in daily diaries. For each sample, the most recent weight from that 265 individual prior to sample collection was included in the analysis. We excluded data 266 from individuals who had recently undergone any potentially stressful procedures (e.g. 267 medical treatment, moves to new enclosures, etc.). The final data sets included 214 268 saliva samples from 36 individuals (mean number of samples per individual = $5.94 \pm$ 269 3.76 SD), and 227 fecal samples from 27 individuals (mean number of samples per 270 individual = 8.41 ± 3.33 SD).

Age (in years) and weight (in g) were included as continuous variables in the analysis. For saliva samples, the mean age of tamarins at the time of sampling was 5.45 years ± 4.00 SD, range 0.21 – 16.43 years; for fecal samples, the mean age was 5.66 years ± 4.75 SD, range 0.72 – 21.88 years. Although wild pied tamarins typically weigh around 430 g (Ford, 1994), weights obtained during this study averaged 501 g + 81 SD, which is comparable to weights obtained from the global captive population of pied tamarins (Species360, 2018).

278 In addition to sex, we also included the following categorical variables:

279	• Rearing history: either hand-reared or parent-reared.
280	• Enclosure type: either on-show (with permanent access to off-exhibit
281	areas) or off-show. All enclosures had both indoor and outdoor areas to
282	which tamarins had permanent access.
283	Social status: dominant or subordinate. Tamarins who were breeding or
284	potentially breeding (i.e. living with a conspecific of the opposite sex)
285	were classed as dominant. Offspring living with their parents were
286	classed as subordinate. Other individuals (i.e. those in single-sex or
287	mixed-species groups) were classified as either dominant or
288	subordinate based on keepers' knowledge of behavior and relationships
289	in each group at the time of sampling (e.g. priority of access to food).
290	• Group composition, categorized as: breeding pair without offspring,
291	family (pair plus offspring), single-sex conspecific group, or mixed-
292	species group (one or two pied tamarins with one or two individuals of
293	another species. Species involved were Leontopithecus chrysopygus, L.
294	chrysomelas, L. rosalia and Callithrix geoffroyi.)
295	• For females in breeding situations, we counted back from the birth of
296	infants to the date each sample was collected. Assuming a gestation
297	period of 160 days (Heistermann, Pröve, Wolters, & Mika, 1987), we
298	classed early pregnancy as 1–80 days gestation and late pregnancy as
299	81–160 days gestation.
200	The dependent variable, cortical level (in pr/ml for calive camples, and pr/EQul

The dependent variable, cortisol level (in ng/ml for saliva samples, and pg/50μl
for fecal samples), was not normally distributed in either case and was therefore

302	natural log-transformed before analysis. We examined the transformed data and
303	standardized residuals graphically to test the assumptions of normality and
304	homoscedasticity, and removed outliers. Final sample sizes for each categorical
305	variable are given in Table 1. We used separate generalized linear mixed models
306	(GLMMs) for each sample type, including individual as a random factor to control for
307	the fact that several samples were obtained from many of the tamarins. Data were
308	analyzed using the statistical software R (version 3.5.1; R Core Development Team,
309	2018) and the packages nlme (Pinheiro, Bates, DebRoy, Sarkar, & R Core Team, 2018),
310	Ime4 (Bates, Maechler, Bolker & Walker, 2015) and MuMIn (Barton, 2015).
311	We first tested for an effect of pregnancy in breeding and potentially breeding
312	females. We examined log-transformed cortisol levels using a GLMM for each sample
313	type, with individual as a random factor, and pregnancy status (early pregnancy, late
314	pregnancy, or not pregnant) as a fixed factor, and adjusted the data sets for
315	subsequent analysis if necessary (see Results).
316	We then ran separate GLMMs for each sample type, including main effects plus an
317	interaction between sex and rearing history, with individual as a random factor. We
318	followed the method outlined by Grueber, Nakagawa, Laws, & Jamieson (2011) for
319	model averaging of GLMMs, and considered models with $\Delta AIC_c < 2$ as having strong
320	support, and those with a ΔAIC_c of >2 to have less support (Burnham and Anderson,
321	2002). We constructed a standardized global model containing all factors of interest,
322	and then used the dredge function in MuMIn to obtain a list of all models with a ΔAIC_c
323	<2 from the best model. We then used model averaging to obtain estimates and 95%

324 confidence intervals for each factor. Finally, means were back-transformed to display325 graphically. The figures were produced in Microsoft Excel.

326

- 327 Ethical statement
- 328 We confirm that the methods used in this study conformed to the UK's Animals
- 329 (Scientific Procedures) Act 1986 Amendment Regulations (SI 2012/3039), the American
- 330 Society of Primatology's Principles for the Ethical Treatment of Non-human Primates,
- and to the Animal Welfare (Jersey) Law 2004, and met the requirements of Durrell
- 332 Wildlife Conservation Trust's Ethics Committee.
- 333 Conflict of interest statement
- 334 The authors confirm that they have no conflicts of interest to declare.
- 335 Results
- 336 Salivary cortisol

We found no effect of pregnancy on cortisol levels in saliva, and so we used all female samples in subsequent analyses. The final GLMM for log cortisol levels in saliva included only two fixed factors, enclosure type and sex; no other model had a ΔAIC_c <2. The influence of enclosure type was in the opposite direction to the predicted one: individuals housed in off-show enclosures had higher salivary cortisol levels than tamarins in on-show exhibits ($\beta = -0.4399 \pm 0.1118$ SE, 95% CI = -0.6887, -0.2196; Figure 1A). Female tamarins had higher salivary cortisol levels than males (β = -0.2905 ± 0.1122 SE, 95% CI = -0.5177, -0.0566; Figure 1B).

345 Fecal cortisol

Although we had few samples from females in late pregnancy, we found some effect of stage of pregnancy on log cortisol levels in fecal samples (F_{2,55} = 2.684, P = 0.077; Figure 2). Examination of paired contrasts showed that females in late pregnancy had significantly higher cortisol levels than those in early pregnancy (P = 0.033), but the other two pairs did not differ significantly. For further analysis, we removed samples from females in late pregnancy (n = 3 samples from two females) from the data set.

353 The MuMIn dredge function produced a set of 12 models with a $\Delta AIC_c < 2$ from the 354 best model (Table 2). The averaged model (Table 3) showed that fecal cortisol levels 355 tended to be lower in subordinate individuals (Figure 3A). There was an interaction 356 between sex and rearing history: hand-reared tamarins of both sexes had higher fecal 357 cortisol levels than parent-reared individuals, but while male parent-reared tamarins 358 had higher cortisol levels than females who were parent-reared, the reverse was true 359 for hand-reared individuals (Figure 3B). Tamarins in off-show enclosures had higher 360 levels than those living in on-show exhibits (Figure 3C), and enclosure type was the 361 only predictor to occur in all 12 models (see Table 2). There was little effect of group 362 composition on cortisol levels (Figure 3D), and this predictor occurred in only one of 363 the averaged models (Table 2). Finally, as individual weight increased, cortisol levels 364 decreased (Figure 4).

365 Discussion

366 As pressure grows on wild populations, captive management will play an 367 increasing role in saving species from extinction, both in terms of maintaining captive 368 populations as insurance and a source for reintroduction, and also in circumstances 369 which require rescue, temporary captivity and translocation to protected 370 environments (Griffiths & Pavajeau, 2008; Baker, Lacy, Leus, & Traylor-Holzer, 2011; 371 Traylor-Holzer, Leus, & Byers, 2018). Understanding the implications of housing, 372 management, hand-rearing and other factors for the wellbeing of these animals is 373 therefore crucial to conservation success (Dickens, Delehanty, & Romero, 2010). In 374 order to obtain robust data, large samples are needed, but this is often difficult in zoo 375 settings as, typically, collections hold only one or two groups of any given species, and 376 cross-institutional studies must incorporate many additional variables to account for 377 differences in the way in which the animals are housed and managed (e.g. 378 Shepherdson et al., 2013). We were able to study a large number of tamarins in a 379 single collection, and therefore draw more reliable conclusions. 380 We found several factors that affected cortisol levels in zoo-housed pied tamarins. 381 However, the results were somewhat different depending on whether cortisol was 382 measured in saliva or feces. Only sex and enclosure type had an effect on salivary 383 cortisol in pied tamarins, while sex, pregnancy, rearing history, social status, weight 384 and enclosure type all influenced fecal cortisol levels. Fecal cortisol may give a better 385 picture of baseline cortisol levels as it is less affected by daily acute events and as such 386 represents the effects of underlying individual, social and environmental factors

387	affecting cortisol production (Millspaugh & Washburn, 2004; Heistermann, 2010). It is
388	also the easiest type of sample to collect for analysis. As salivary cortisol represents an
389	individual's immediate reaction to an event (Kuhar, Bettinger, & Laudenslager, 2005;
390	Laudenslager, Bettinger, & Sackett, 2006), it may therefore be of most help in
391	investigating the impact of acute stressors on the wellbeing of pied tamarins.
392	Unfortunately we were not able to make direct comparisons between fecal and
393	salivary cortisol levels in individuals, as the two sets of samples were collected at
394	different times.
395	It is also possible that, since samples from the two media were collected at
396	different times of the year (saliva: January–June, feces: May–November), cortisol
397	levels were differentially affected by weather conditions – both temperature and
398	precipitation have been shown to influence cortisol (de Bruijn and Romero, 2018).
399	Finally, previous research suggests that salivary and fecal cortisol may manifest
400	different response patterns to stress. For example, most empirical research with
401	callitrichids has found increased fecal cortisol following a stressor (e.g. Galvão-Coelho,
402	Silva, & De Sousa, 2012; Pizzuto et al., 2015), whereas recent studies in the common
403	marmoset Callithrix jacchus have reported that salivary cortisol following exposure to a
404	stressor may either decrease (Ash et al., 2018; Cross and Rogers, 2006) or increase
405	(Kaplan et al., 2012).

406

407 Individual factors affecting cortisol levels

408 We found no effect of age on cortisol in either feces or saliva. Prediction 1, that 409 older pied tamarins would exhibit higher cortisol levels as they may be more 410 susceptible to wasting syndrome (Smithyman, 2012), was therefore not supported. 411 Similar results have been reported by Bales et al. (2006) for fecal cortisol in male 412 golden lion tamarins, *Leontopithecus rosalia*. Studies in other primate taxa have 413 produced inconsistent results: for example, Erwin, Tigno, Gerzanich, & Hansen (2004) 414 found a positive correlation between age and plasma cortisol in *M. mulatta*, while in 415 the same species, Dettmer, Novak, Suomi, & Meyer (2012) reported a decrease in hair 416 cortisol with age, and Fourie et al. (2015a) found that cortisol levels in two baboon 417 species (Papio anubis, P. hamadryas) were lowest in adulthood, and higher in both 418 young and older age groups. It is likely that other factors, such as the species' social 419 system (Abbott et al., 2003), housing conditions, and individual reproductive status, 420 have a greater influence on cortisol levels than age. 421 Weight loss is one of the most consistent and pronounced changes during 422 exposure to stress (Tamashiro et al., 2005). We found no effect of weight on salivary 423 cortisol, but fecal cortisol decreased significantly as weight increased, as we expected

424 (Prediction 2). Links have been found in a number of taxa between poorer body

425 condition or lower weight, and higher baseline cortisol levels (e.g. Macbeth, Cattet,

426 Obbard, Middel, & Janz, 2012; Cattet et al., 2014; Trevisan et al., 2017) or higher

427 glucocorticoid reactivity (Breuner & Hahn, 2003; Pereyra & Wingfield, 2003). Changes

428 in weight may therefore give an early indication of both increased stress levels and the

429 possibility of wasting syndrome, and thus regular non-disruptive weighing, as used in 430 our colony, is an important tool in monitoring health and wellbeing in tamarins in 431 captivity. We excluded data from tamarins with symptoms of illness, but it is possible 432 that individual health parameters could have confounded cortisol titers in our subjects. 433 Pregnancy had no effect on cortisol levels in saliva, but as expected, cortisol in 434 feces was highest in females in late pregnancy (Prediction 3). Although our sample size 435 for females in late pregnancy was small, this is consistent with the typical pattern in 436 primates, including humans (Ziegler et al., 1995; Leung et al., 2001; Bales et al., 2005). 437 It is well known that the reproductive status of the female in addition to pregnancy 438 influences cortisol titers, e.g. lactation (Starling, Charpentier, Fitzpatrick, Scordato, & 439 Drea, 2010). In the closely related common marmoset, *Callithrix jacchus*, cortisol 440 levels vary reliably across the ovarian cycle and are significantly raised around the peri-441 ovulatory phase (Saltzman, Schultz-Darken, Wegner, Wittwer, & Abbott, 1998). In our 442 study we were only able to control for early and late pregnancy, and did not take into 443 account other reproductive phases such as ovulation that could have influenced 444 cortisol titers. Furthermore, in several primate species, baseline cortisol levels of 445 males are also affected by the reproductive status of the females in the group, in 446 particular ovulation – something that we could not take into account when analyzing 447 male cortisol values but which may have affected our data (Surbeck, Deschner, 448 Weltring, & Hohmann, 2012; Schoof, Jack & Ziegler, 2014). 449 We found that female pied tamarins had higher salivary cortisol levels than males 450 (Prediction 4). This agrees with a previous study of four zoo-housed pied tamarins that 451 found significantly higher fecal cortisol levels in the two females (Armstrong &

452 Santymire, 2012), and with studies of urinary and fecal cortisol in other callitrichid
453 species (Smith & French, 1997a; Wark et al. 2016).

454 The effect of sex on cortisol in feces in our study, however, depended on the 455 rearing history of the individual. Overall, hand-reared tamarins had higher fecal 456 cortisol levels than parent-reared individuals, supporting Prediction 5, but while hand-457 reared females did have higher cortisol levels than hand-reared males, male parent-458 reared tamarins had higher cortisol levels than females who were parent-reared 459 (Predictions 4 and 5). This contrasts with the results reported for Saguinus geoffroyi 460 by Kuhar, Bettinger, Sironen, Shaw, & Lasley (2003), who found no differences in 461 cortisol between hand-reared and parent-reared individuals. 462 It is important to note that the number of hand-reared individuals included in our 463 study was small (n = 5). However, the generally higher fecal cortisol levels that we 464 found in hand-reared pied tamarins are consistent with the greater incidence of 465 negative behavior towards humans that we have observed in hand-reared tamarins 466 (Coe, 2014). Similar results have been reported for pileated gibbons, Hylobates 467 pileatus, by Pirovino et al. (2011): hand-reared gibbons had higher levels of fecal 468 glucocorticoid metabolites and exhibited more abnormal behavior than parent-reared 469 individuals. Hand- or nursery-rearing of primates is known to affect allostatic load (a 470 composite measure of stress; Edes, Wolfe, & Crews, 2016), brain development (Bogart, 471 Bennett, Schapiro, Reamer, & Hopkins, 2014), behavior, including parenting (Mallapur 472 & Choudhury, 2003; Niebruegge & Porton, 2006; Vermeer & Devreese, 2015) and 473 response to stressors in later life (Dettmer et al., 2012), and in chimpanzees, the higher

474 the level of conspecific as opposed to human interaction throughout life, the lower the 475 level of cortisol in hair samples (Jacobson, Freeman, Santymire & Ross, 2018). Hand-reared tamarins are often poor parents, probably in part because they are 476 477 less likely to have had experience caring for infants as helpers in family groups (Tardif, 478 Richter & Carson, 1984; but see Baker & Woods, 1992, for an exception). Interestingly, 479 while hand-reared female pied tamarins in our colony have invariably been 480 incompetent parents (Price et al., 2016) we have documented adequate parental care 481 in one hand-reared male. Although our sample is too small to draw any firm 482 conclusions, the relationships between rearing history, cortisol levels and parenting, 483 and how they may differ between the sexes, will be important to investigate further. 484 The possibility that these relationships may differ from one species to another should 485 also be borne in mind. 486 Our results also raise questions about the role and ethics of hand-rearing pied 487 tamarins. Attitudes towards hand-rearing primates in zoos have changed since the 488 1950s, when it was frequently practiced in order to increase survival rates (Porton & 489 Niebruegge, 2006); realization of the importance of parental rearing, in particular for 490 social development, has led to a decrease in the number of primates removed by zoos 491 for hand-rearing and an improved understanding of the need for early socialization 492 with conspecifics. However, for threatened species such as pied tamarins, hand rearing 493 has remained an important tool to increase captive population sizes. Although in 494 modern zoos, hand-reared infants are socialized from a very early age, work remains 495 to be done in developing rearing and socialization methods that result in adults that 496 are behaviorally and physiologically indistinguishable from parent-reared individuals.

497

498 Social factors affecting cortisol levels

499 We found no effect of social status on salivary cortisol, but there was a trend for 500 subordinate tamarins to have lower fecal cortisol levels than tamarins classed as 501 dominant. These results supports Prediction 6 and are in agreement with previous 502 studies of a number of callitrichid species using various methods of measuring cortisol 503 levels, all of which have found either no effect of status, or reduced cortisol levels in 504 subordinate individuals (Baker, Abbott & Saltzman, 1999; Saltzman et al. 2004: plasma 505 cortisol in captive Callithrix jacchus; Smith & French, 1997b: urinary cortisol in C. kuhli; 506 Huck, Löttker, Heymann, & Heistermann, 2005: fecal cortisol in wild Saguinus mystax; 507 Bales et al., 2005, 2006: fecal cortisol in wild *Leontopithecus rosalia*). In a synthesis of 508 published data on cortisol and status in primates in relation to social systems, Abbott 509 et al. (2003) pointed out that the frequency and rate at which subordinates receive 510 aggression, the stability of access to resources and social relationships, and coping 511 strategies, may all be as important as low rank itself in mediating the stress response. 512 In callitrichids, social life is characterized by cooperation, usually among close relatives, 513 and a high level of affiliative interactions (Schaffner & Caine, 2000), and thus 514 subordinates are not subject to frequent stressors (Abbott et al., 2003). Our results 515 from pied tamarins support this view. However, in some species, rank-related patterns 516 of glucocorticoid excretion are only evident under certain conditions, such as during 517 times of food shortages (L. catta; Cavigelli, 1999), the mating season (e.g. Brachyteles 518 arachnoides hypoxanthus; Strier, Lynch & Ziegler, 2003), pregnancy (semi-free-ranging

519 provisioned L. catta; Starling et al., 2010) or lactation (Cercopithecus mitis; Foerster, 520 Cords & Monfort, 2011). Our study might have yielded different relationships between 521 cortisol and social rank under different social or environmental conditions such as 522 following a birth or aggression. 523 in contrast to other primate studies in which variations in group composition such 524 as group size (Pride, 2005) or number of males in a group (Smith, McCusker, Stevens & 525 Elwood, 2015) have been shown to have a significant impact on cortisol levels, we 526 found little influence of group composition on cortisol levels in either saliva or feces. 527 Our expectation that pied tamarins would exhibit higher cortisol levels if they were 528 housed with members of other species was therefore not supported (Prediction 7). 529 This suggests that as long as an individual is housed socially, it does not make a great 530 deal of difference whether it is living in a mixed-sex family or pair, or a single-sex 531 group, or even with individuals from another species, and is encouraging in terms of 532 management strategies for captive tamarins. However, it is important to remember 533 that each situation is different, and any new grouping, whether of conspecifics or not, 534 should be closely monitored (Buchanan-Smith, 2012).

535

536 Environmental factors affecting cortisol levels

537 We found that in both saliva and feces, cortisol levels were significantly higher in 538 tamarins living in enclosures that were not on show to the public (Prediction 8). This 539 was unexpected – the presence of visitors has often been highlighted as a source of 540 stress and a cause of abnormal behavior in zoo primates (Hosey, 2000), and an earlier

541 study (Armstrong & Santymire, 2012) found that a pair of pied tamarins on exhibit had 542 higher levels of fecal glucocorticoid metabolites than a pair living off-show. However, 543 our results are in agreement with previous behavioral studies of pied tamarins at 544 Jersey Zoo, which found that visitors had no effect on levels of stress-related behaviors 545 (Holm et al., 2012), and that tamarins housed on-show were less vigilant, and vocalized 546 and scent-marked less often than tamarins in off-show enclosures (Steinbrecher, 547 2016). We suggest that this may be because tamarins on show to the public at the zoo 548 have high and relatively naturalistic enclosures with stand-off barriers preventing 549 direct contact with the public (Figure 5), and they are also able to retreat indoors 550 whenever they wish. Off-show enclosures are similar in design, but typically house 551 more groups of pied tamarins per building (though fewer groups overall) than the on-552 show enclosures. Crowding has been linked to elevated cortisol in several primate 553 species and may have contributed to the raised levels in the tamarins off-show 554 (Dettmer, Novak, Meyer, & Suomi, 2014; Gabriel, Gould, & Cook, 2018; Pearson, 555 Reeder & Judge, 2015). Interestingly, Pirovino et al. (2011) also found that cortisol 556 values varied across pileated gibbons not only living in different institutions, but across 557 animals residing in different enclosures within the same institution, illustrating the 558 mixture of factors that may affect cortisol at a local level.

559 Cabana et al. (2018) reported that the likelihood of wasting syndrome developing 560 in zoo-housed callitrichids was lower if there were visual barriers between visitors and 561 monkeys, and if the animals had safe areas to which they could retreat. Another study 562 at Jersey Zoo demonstrated that the level of noise in a building (mostly due to other 563 animals housed there) was positively related to cortisol levels in *S. bicolor* (Simpkins,

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564 Routh, Wormell, & Price, 2013). Similarly, Kuhar et al. (2003) found that colony 565 housed S. geoffroyi showed higher levels of aggression and lower activity levels than 566 non-colony-housed tamarins, and higher levels of physical activity have been shown to 567 mitigate stress responses in women (Puterman et al. 2011). It would therefore be 568 interesting to investigate activity levels in tamarins housed on- and offshow, and to 569 monitor changes in both behavior and cortisol levels in groups that are moved from 570 one enclosure type to another. Further research into how enclosure design, 571 management, housing density and proximity to conspecifics versus other species 572 influence behavior and stress levels in pied tamarins and other callitrichids is therefore 573 needed. 574 Although our sample size was comparatively large, we did not have sufficient data 575 to be able to take into account other non-stress related and stress-related factors that

577 Reale, 2008; Shepherdson et al., 2013), which influences cortisol in another callitrichid

might affect HPA function and thus cortisol levels, such as personality (Martin and

578 species, the common marmoset (Inoue-Murayama et al., 2018). Variations in

579 metabolic rate (Goymann 2012), activity levels (Smith, McGreer-Whitworth, & French,

580 1998) and amount of keeper interaction (Carlstead, Paris, & Brown, 2018) were not

581 controlled in this study, but may also affect cortisol levels. Finally, seasonal variation in

temperature or poor weather (which could reduce visitor numbers, but potentially

583 increase intragroup tension as tamarins stayed indoors for longer periods) may also

have had an effect on cortisol in this study. For example, it would be interesting to

585 investigate levels of salivary cortisol when tamarins are indoors versus outdoors.

586 Focused studies on these possibilities, and further research into the factors we

587 investigated, would contribute a great deal to the successful management of this and

588 other threatened callitrichid species both in captivity, and in cases where intensive

589 management of free-living populations becomes necessary.

590

591 Conclusions

592	1.	Fewer factors affect cortisol in saliva than in feces in captive pied tamarins;
593		salivary cortisol is therefore likely to be of more value in assessing the
594		immediate impact of stressful events than in understanding underlying
595		sources of chronic stress.
596	2.	Female pied tamarins have higher fecal cortisol levels in late pregnancy.
597	3.	Female tamarins have higher salivary cortisol levels than males.
598	4.	Female hand-reared tamarins have higher fecal cortisol levels than males,
599		but the reverse is true for parent-reared tamarins.
600	5.	Hand-reared pied tamarins have higher levels of fecal cortisol overall and
601		this is consistent with the higher levels of abnormal and aggressive
602		behavior seen in hand-reared tamarins.
603	6.	Fecal cortisol levels increase in pied tamarins as weight decreases, which
604		can be an indicator of wasting syndrome in this species. Regular non-
605		disruptive weighing may help to identify tamarins at risk.
606	7.	Pied tamarins housed on show had lower cortisol levels in both saliva and
607		feces than tamarins living in off-show enclosures with a higher number of
608		conspecifics in a given area, and therefore may not be affected by

609	disturbance from visitors as long as their contact with people is minimized
610	by the use of barriers and access to off-exhibit areas.
611	8. Age does not affect levels of cortisol in this species.
612	Acknowledgements
613	We thank Fiona Gladwell at the Central Science Laboratory for analysis of saliva
614	samples; Philippa Blanco, Will Masefield and the many other members of the Mammal
615	Department at Jersey Zoo who helped to collect samples; Valerie Harding for
616	assistance in compiling data from daily diaries; and Lianne Concannon, Tine Griede,
617	and Henry Kuijpers for advice on the statistical analyses. We are also grateful to Karen
618	Bales and two anonymous reviewers for constructive comments on previous drafts.
619	
620	References
621	Abbott, D. H., Keverne, E. B., Bercovitch, F. B., Shively, C. A., Mendoza, S. P., Saltzman,
622	W., Snowdon, C. T., Ziegler, T. E., Banjevic, M., Garland Jr, T. & Sapolsky, R. M.,
623	(2003). Are subordinates always stressed? A comparative analysis of rank
624	differences in cortisol levels among primates. Hormones and Behavior, 43, 67–

625 82. https://doi.org/10.1016/s0018-506x(02)00037-5

Anzenberger, G. & Falk, B. (2012). Monogamy and family life in callitrichid monkeys:
deviations, social dynamics and captive management. *International Zoo Yearbook,* 46, 109–122. https://doi.org/10.1111/j.1748-1090.2012.00176.x

629	Armstrong, D. & Santymire, R. (2013). Hormonal and behavioral variation in pied
630	tamarins housed in different management conditions. Zoo Biology, 32, 299–306.
631	https://doi.org/10.1002/zoo.21023
632	Ash, H., Smith, T. E., Knight, S. & Buchanan-Smith, H. M. (2018). Measuring
633	physiological stress in the common marmoset (Callithrix jacchus): Validation of a
634	salivary cortisol collection and assay technique. Physiology and Behavior, 185,
635	14–22. https://doi.org/10.1016/j.physbeh.2017.12.018
636	Baker, A. J. & Woods, F. (1992). Reproduction of the emperor tamarin (Saguinus
637	<i>imperator</i>) in captivity, with comparisons to cotton-top and golden lion tamarins.
638	American Journal of Primatology, 26, 1–10.
639	https://doi.org/10.1002/ajp.1350260104
640	Baker, A. M., Lacy, R. C., Leus, K. & Traylor-Holzer, K. (2011). Intensive management of
641	populations for conservation. WAZA Magazine: Towards Sustainable Population
642	Management, 12, 40–43.
643	Baker, J. V., Abbott, D. H. & Saltzman, W. (1999). Social determinants of reproductive
644	failure in male common marmosets housed with their natal family. Animal
645	<i>Behaviour</i> , 58, 501–513. https://doi.org/10.1006/anbe.1999.1200
646	Bales, K., French, J., Hostetler, C. & Dietz, J. (2005). Social and reproductive factors
647	affecting cortisol levels in wild female golden lion tamarins (Leontopithecus
648	rosalia). American Journal of Primatology, 67, 25–35.
649	https://doi.org/10.1002/ajp.20167

- Bales, K. L., French, J. A., McWilliams, J., Lake, R. A. & Dietz, J. M. (2006). Effects of
- 651 social status, age, and season on androgen and cortisol levels in wild male golden
- lion tamarins (*Leontopithecus rosalia*). *Hormones and Behavior*, 49, 88–95.
- 653 https://doi.org/10.1016/j.yhbeh.2005.05.006
- Barton, K. (2015). *MuMIn: Multi-model inference*. R package (Version 1.13.4).
- 655 http://CRAN.Rproject.org/package=MuMIn
- 656 Bates, D., Maechler, M., Bolker, B., & Walker, S. (2014). Ime4: Linear mixed-effects
- 657 models using Eigen and S4. R package version 1.1-7, http://CRAN.R-
- 658 project.org/package=lme4.
- Beck, B. B. & Power, M. L. (1988). Correlates of sexual and maternal competence in
 captive gorillas. *Zoo Biology*, 7, 339–350.
- 661 https://doi.org/10.1002/zoo.1430070405
- 662 Beehner, J. C. & Bergman, T. J. (2017). The next step for stress research in primates: to
- 663 identify relationships between glucocorticoid secretion and fitness. *Hormones*
- 664 *and Behavior* 91, 68–83. <u>https://doi.org/10.1016/j.yhbeh.2017.03.003</u>
- 665 Behie, A.M., Pavelka, M.S. and Chapman, C.A., 2010. Sources of variation in fecal
- 666 cortisol levels in howler monkeys in Belize. *American Journal of Primatology*, 72,
 667 600-606.
- 668 Bergman, T. J., Beehner, J. C., Cheney, D. L., Seyfarth, R. M. & Whitten, P. L. (2005).
- 669 Correlates of stress in free-ranging male chacma baboons, *Papio hamadryas*

- 670 *ursinus*. *Animal Behaviour*, 70, 703–713.
- 671 https://doi.org/10.1016/j.anbehav.2004.12.017
- 672 Bogart, S. L., Bennett, A. J., Schapiro, S. J., Reamer, L. A. & Hopkins, W. D. (2014).
- 673 Different early rearing experiences have long-term effects on cortical
- 674 organization in captive chimpanzees (Pan troglodytes). Developmental Science,
- 675 17, 161–174. https://doi.org/10.1111/desc.12106
- 676 Breuner, C. W. & Hahn, T. P. (2003). Integrating stress physiology, environmental
- 677 change, and behavior in free-living sparrows. Hormones and Behavior, 43, 115–
- 678 123. https://doi.org/10.1016/s0018-506x(02)00020-x
- de Bruijn, R. & Romero, L.M. (2018). The role of glucocorticoids in the vertebrate
- 680 response to weather. *General and Comparative Endocrinology*, 269, 1–23.
- 681 https://doi.org/10.1016/j.ygcen.2018.07.007
- 682 Buchanan-Smith, H. M. (2012). Mixed-species exhibition of neotropical primates:
- 683 analysis of species combination success. International Zoo Yearbook, 46, 150–
- 684 163. https://doi.org/10.1111/j.1748-1090.2011.00151.x
- 685 Burnham, K. P. & Anderson, D. R. (2002). Model Selection and Multimodel Inference: A
- 686 *Practical Information-Theoretic Approach*. New York: Springer.
- 687 https://doi.org/10.1007/b97636

688	Cabana, F., Maguire, R., Hsu, CD. & Plowman, A. (2018). Identification of possible
689	nutritional and stress risk factors in the development of marmoset wasting
690	syndrome. Zoo Biology 37, 98–106. https://doi.org/10.1002/zoo.21398
691	Carlstead, K. & Brown, J. L. (2005). Relationships between patterns of fecal corticoid
692	excretion and behavior, reproduction, and environmental factors in captive black
693	(Diceros bicornis) and white (Ceratotherium simum) rhinoceros. Zoo Biology, 24,
694	215–232. https://doi.org/10.1002/zoo.20050
695	Carlstead, K., Brown, J. L. & Seidensticker, J. (1993). Behavioral and adrenocortical
696	responses to environmental changes in leopard cats (Felis bengalensis). Zoo
697	<i>Biology</i> , 12, 321–331. https://doi.org/10.1002/zoo.1430120403
698	Carlstead, K., Paris, S. & Brown, J. L. (2018). Good keeper-elephant relationships in
699	North American zoos are mutually beneficial to welfare. Applied Animal
700	Behaviour Science, 211, 103–111.
701	https://doi.org/10.1016/j.applanim.2018.11.003
702	Cattet, M., Macbeth, B. J., Janz, D. M., Zedrosser, A., Swenson, J. E., Dumond, M., &
703	Stenhouse, G. B. (2014). Quantifying long-term stress in brown bears with the
704	hair cortisol concentration: a biomarker that may be confounded by rapid
705	changes in response to capture and handling, <i>Conservation Physiology</i> , 2, 1–15.
706	https://doi.org/10.1093/conphys/cou026

707	Cavigelli, S. A. (1999). Behavioural patterns associated with faecal cortisol levels in
708	free-ranging female ring-tailed lemurs, Lemur catta. Animal Behaviour, 57, 935–
709	944. <u>https://doi.org/10.1006/anbe.1998.1054</u>
710	Cerda-Molina, A.L., Hernández-López, L., Díaz-Díaz, G., Mejía-Varas, F., Chavira, R. and
711	Mondragón-Ceballos, R. (2012). Housing with females increases testosterone and
712	cortisol levels in captive groups of black-handed spider monkeys (Ateles
713	geoffroyi). Zoo Biology, 31, 490-497.
714	Clark, F. E., Fitzpatrick, M., Hartley, A., King, A. J., Lee, T., Routh, A., Walker, S. L. &
715	George, K. (2012). Relationship between behavior, adrenal activity, and
716	environment in zoo-housed western lowland gorillas (Gorilla gorilla gorilla). Zoo
717	<i>Biology</i> , 31, 306–321. https://doi.org/10.1002/zoo.20396
718	Clubb, R., Rowcliffe, M., Lee, P., Mar, K. U., Moss, C. & Mason, G. J. (2008).
719	Compromised survivorship in zoo elephants. Science, 322, 1649–1649.
720	https://doi.org/10.1126/science.1164298
721	Coe, A. B. (2014). Hand-reared vs parent-reared: reproductive success and behaviour
722	of pied tamarins (Saguinus bicolor) housed at Durrell Wildlife Conservation Trust.
723	Nottingham Trent University, UK (unpublished BSc dissertation).
724	Cook, N. J. (2012). Minimally invasive sampling media and the measurement of
725	corticosteroids as biomarkers of stress in animals. Canadian Journal of Animal
726	Science, 92, 227–259. https://doi.org/10.4141/cjas2012-045

727	Cross, N., Pines, M. K. & Rogers, L. J. (2004). Saliva sampling to assess cortisol levels in
728	unrestrained common marmosets and the effect of behavioral stress. American
729	Journal of Primatology, 62, 107–114. https://doi.org/10.1002/ajp.20005
730	Cross, N., & Rogers, L. J. (2004). Diurnal cycle in salivary cortisol levels in common
731	marmosets. Developmental Psychobiology, 45, 134–139.
732	https://doi.org/10.1002/dev.20023
733	Cross, N. & Rogers, L.J. (2006). Mobbing vocalisations as a coping response in the
734	common marmoset. Hormones and Behaviour, 49, 237–245.
735	https://doi.org/10.1016/j.yhbeh.2005.07.007
736	Dawkins, M. (2004). Using behaviour to assess animal welfare. Animal Welfare, 13, S3-
737	7.
738	Dettmer, A. M., Novak, M. A., Suomi, S. J. & Meyer, J. S. (2012). Physiological and
739	behavioral adaptation to relocation stress in differentially reared rhesus
740	monkeys: hair cortisol as a biomarker for anxiety-related responses.
741	Psychoneuroendocrinology, 37, 191–199.
742	https://doi.org/10.1016/j.psyneuen.2011.06.003
743	Dettmer, A. M., Novak, M. A., Meyer, J. S. & Suomi, S. J. (2014). Population density-
744	dependent hair cortisol concentrations in rhesus monkeys (Macaca mulatta).
745	Psychoneuroendocrinology, 42, 59–67.
746	https://doi.org/10.1016/j.psyneuen.2014.01.002

747	Diamandis, E. P. & Christopoulos, T. K., Eds. (1996). Immunoassay. New York: Academic
748	Press. https://doi.org/10.1016/b978-0-12-214730-2.x5000-6
749	Dickens, M. J., Delehanty, D. J. & Romero, L. M. (2010). Stress: an inevitable
750	component of animal translocation. Biological Conservation, 143, 1329–1341.
751	https://doi.org/10.1016/j.biocon.2010.02.032
752	EAZA (2013). Strategic Plan 2013–2016. Amsterdam: EAZA.
753	https://www.eaza.net/assets/Uploads/Strategies/2013-2016-EAZA-Strategy-low-
754	<u>res-version2.pdf</u>
755	Edes, A. N., Wolfe, B. A. & Crews, D. E. (2016). Rearing history and allostatic load in
756	adult western lowland gorillas (Gorilla gorilla gorilla) in human care. Zoo Biology,
757	35, 167–173. https://doi.org/10.1002/zoo.21270
758	Erwin, J. M., Tigno, X. T., Gerzanich, G. & Hansen, B. C. (2004). Age-related changes in
759	fasting plasma cortisol in rhesus monkeys: implications of individual differences
760	for pathological consequences. The Journals of Gerontology Series A: Biological
761	Sciences and Medical Sciences, 59, B424–B432.
762	https://doi.org/10.1093/gerona/59.5.b424
763	Foerster, S., Cords, M. & Monfort, S.L. (2011). Social behavior, foraging strategies, and
764	fecal glucocorticoids in female blue monkeys (Cercopithecus mitis): potential
765	fitness benefits of high rank in a forest guenon. American Journal of

766 *Primatology*, 73, 870–882. https://doi.org/10.1002/ajp.20955

- Ford, S. M. (1994). Evolution of sexual dimorphism in body weight in platyrrhines.
- 768 American Journal of Primatology, 34, 221–244.
- 769 https://doi.org/10.1002/ajp.1350340211
- 770 Fourie, N. H., Jolly, C. J., Phillips-Conroy, J. E., Brown, J. L. & Bernstein, R. M. (2015a).
- 771 Variation of hair cortisol concentrations among wild populations of two baboon
- species (*Papio anubis*, *P. hamadryas*) and a population of their natural hybrids.
- 773 *Primates*, 56, 259–272. https://doi.org/10.1007/s10329-015-0469-z
- Fourie, N. H., Turner, T. R., Brown, J. L., Pampush, J. D., Lorenz, J. G. & Bernstein, R. M.
- 775 (2015b). Variation in vervet (*Chlorocebus aethiops*) hair cortisol concentrations
- reflects ecological disturbance by humans. *Primates*, 56, 365–373.
- 777 https://doi.org/10.1007/s10329-015-0486-y
- 778 Gabriel, D. N., Gould, L. & Cook, S. (2018). Crowding as a primary source of stress in an
- endangered fragment-dwelling strepsirrhine primate. Animal Conservation, 21,
- 780 76–85. https://doi.org/10.1111/acv.12375
- 781 Galvão-Coelho, N. L., Silva, H. P. A. and De Sousa, M. B. C. (2012). The influence of sex
- and relatedness on stress response in common marmosets (Callithrix
- *jacchus*). *American Journal of Primatology*, 74, 819–827.
- 784 https://doi.org/10.1002/ajp.22032
- 785 Garber, P. A. (1997). One for all and breeding for one: Cooperation and competition as
- a tamarin reproductive strategy. *Evolutionary Anthropology*, 5, 187–199.

787 https://doi.org/10.1002/(sici)1520-6505(1997)5:6%3C187::aid-

788 evan1%3E3.0.co;2-a

789 Gentry, L. & Margulis, S. W. (2008). Behavioral effects of introducing pied tamarin

790 (Saguinus bicolor) to black howler monkey (Alouatta caraya) and white-faced

791 saki (Pithecia pithecia) in a zoological park. American Journal of Primatology, 70,

792 1–5. https://doi.org/10.1002/ajp.20521

793 Gladwell, F. & Pick, J. (2007). Salivary cortisol in pied tamarins (Saguinus bicolor): Non-

794 invasive measure of stress. Poster presented at the 6th World Congress on

795 Stress, 11–13 October 2007, Vienna, Austria.

Goldizen, A. W. (1987). Facultative polyandry and the role of infant-carrying in wild
 saddle-back tamarins (*Saquinus fuscicollis*). *Behavioral Ecology and Sociobiology*,

798 20, 99–109. https://doi.org/10.1007/bf00572631

Gordo, M., Calleia, F. O., Vasconcelos, S. A., Leite, J. J. & Ferrari, S. F. (2013). The

800 challenges of survival in a concrete jungle: conservation of the pied tamarin

801 (Saguinus bicolor) in the urban landscape of Manaus, Brazil. In L. K. Marsh & C. A.

802 Chapman (Eds.), Primates in fragments: complexity and resilience (pp. 357–370).

803 Springer: New York. https://doi.org/10.1007/978-1-4614-8839-2_23

804 Goymann, W. (2012). On the use of non-invasive hormone research in uncontrolled,

805 natural environments: the problem with sex, diet, metabolic rate and the

806 individual. *Methods in Ecology and Evolution*, 3, 757–765.

807 https://doi.org/10.1111/j.2041-210x.2012.00203.x

808	Griffiths, R. A. & Pavajeau, L. (2008). Captive breeding, reintroduction, and the
809	conservation of amphibians. Conservation Biology, 22, 852–861.
810	https://doi.org/10.1111/j.1523-1739.2008.00967.x
811	Gross, M. (2015). Can zoos offer more than entertainment? Current Biology, 25, R391–
812	R394. https://doi.org/10.1016/j.cub.2015.04.056
813	Grueber, C. E., Nakagawa, S., Laws, R. J. & Jamieson, I.G. (2011). Multimodel inference
814	in ecology and evolution: challenges and solutions. Journal of Evolutionary
815	<i>Biology</i> , 24, 699–711. https://doi.org/10.1111/j.1420-9101.2010.02210.x
816	Hart, K. (2012). The use of cortisol for the objective assessment of stress in animals:
817	Pros and cons. The Veterinary Journal, 192, 137–139.
818	Heistermann, M. (2010). Non-invasive monitoring of endocrine status in laboratory
819	primates: methods, guidelines and applications. Advances in Science and
820	Research, 5, 1–9. https://doi.org/10.1016/j.tvjl.2012.03.016
821	Heistermann, M., Pröve, E., Wolters, H. J. & Mika, G. (1987). Urinary oestrogen and
822	progesterone excretion before and during pregnancy in a pied bare-face tamarin
823	(Saguinus bicolor bicolor). Journal of Reproduction and Fertility, 80, 635–640.
824	https://doi.org/10.1530/jrf.0.0800635
825	Henry, M. D., Hankerson, S. J., Siani, J. M., French, J. A. & Dietz, J. M. (2013). High rates
826	of pregnancy loss by subordinates leads to high reproductive skew in wild golden

- lion tamarins (*Leontopithecus rosalia*). *Hormones and Behavior*, 63, 675–683.
- 828 https://doi.org/10.1016/j.yhbeh.2013.02.009
- 829 Heymann, E. W & Buchanan-Smith, H. M. (2000). The behavioural ecology of mixed-
- 830 species troops of callitrichine primates. *Biological Reviews*, 75, 169–190.
- 831 https://doi.org/10.1111/j.1469-185x.1999.tb00044.x
- 832 Holm, C., Priston, N., Price, E. & Wormell, D. (2012). An investigation into the
- 833 environmental factors affecting behavioural stress in captive pied tamarins,
- 834 Saguinus bicolor. Canopy, 12, 14–15. <u>https://www.brookes.ac.uk/social-</u>
- 835 <u>sciences/courses/primate-conservation/canopy/</u>
- Hosey, G. (2000). Zoo animals and their human audiences: what is the visitor effect?
 Animal Welfare, 9, 343–357.
- 838 Huck, M., Löttker, P., Heymann, E. W. & Heistermann, M. (2005). Characterization and
- 839 social correlates of fecal testosterone and cortisol excretion in wild male
- 840 Saguinus mystax. International Journal of Primatology, 26, 159–179.
- 841 https://doi.org/10.1007/s10764-005-0728-8
- 842 Ialeggio, D. M., & Baker, A. J. (1995). Results of a preliminary survey into wasting
- 843 marmoset syndrome in callitrichid collections. In Proceedings of the First
- 844 Conference on Zoo and Wildlife Nutrition, pp. 148–158.
- 845 ICMBio (2011). Executive Summary of National Action Plan for the Conservation of the
 846 Pied Tamarin. Brasilia: Instituto Chico Mendes. Available at:

- 847 www.icmbio.gov.br/biodiversidade/fauna-brasileira/lista-planos-de-acao-848 nacionais
- 849 Inoue-Murayama, M., Yokoyama, C., Yamanashi, Y. & Weiss, A. (2018). Common
- 850 marmoset (*Callithrix jacchus*) personality, subjective well-being, hair cortisol level
- and AVPR1a, OPRM1, and DAT genotypes. *Scientific Reports*, 8, 10255.
- 852 https://doi.org/10.1038/s41598-018-28112-7
- Jacobson, S. L., Freeman, H. D., Santymire, R. M. & Ross, S. R. (2018). Atypical
- 854 experiences of captive chimpanzees (*Pan troglodytes*) are associated with higher
- hair cortisol concentrations as adults. *Royal Society Open Science*, 4, 170932.
- 856 https://doi.org/10.1098/rsos.170932
- 857 Kagan, R., Carter, S. & Allard, S. (2015). A universal animal welfare framework for zoos.
- 358 Journal of Applied Animal Welfare Science, 18 (suppl.1), S1–S10.
- 859 https://doi.org/10.1080/10888705.2015.1075830
- 860 Kaplan, G., Pines, M. K. & Rogers, L. J. (2012). Stress and stress reduction in common
- 861 marmosets. *Applied Animal Behaviour Science*, 137, 175–182.
- 862 https://doi.org/10.1016/j.applanim.2011.04.011
- 863 Keay, J. M., Singh, J., Gaunt, M. C. & Kaur, T. (2006). Fecal glucocorticoids and their
- 864 metabolites as indicators of stress in various mammalian species: a literature
- 865 review. Journal of Zoo and Wildlife Medicine, 37, 234–244.
- 866 https://doi.org/10.1638/05-050.1

- Keulartz, J. (2015). Captivity for conservation? Zoos at a crossroads. *Journal of Agricultural and Environmental Ethics*, 28, 335–351.
- 869 https://doi.org/10.1007/s10806-015-9537-z
- 870 King, N. E. & Mellen, J. D. (1994). The effects of early experience on adult copulatory
- behavior in zoo-born chimpanzees (*Pan troglodytes*). *Zoo Biology*, 13, 51–59.
- 872 https://doi.org/10.1002/zoo.1430130107
- 873 Kuhar, C. W., Bettinger, T. L. & Laudenslager, M. L. (2005). Salivary cortisol and
- behaviour in an all-male group of western lowland gorillas (*Gorilla g. gorilla*).
- 875 *Animal Welfare*, 14, 187–193.
- 876 Kuhar, C. W., Bettinger, T. L., Sironen, A. L., Shaw, J. H. & Lasley, B. L. (2003). Factors
- 877 affecting reproduction in zoo-housed Geoffroy's tamarins (*Saguinus geoffroyi*).
- 878 Zoo Biology, 22, 545–559. https://doi.org/10.1002/zoo.10099
- 879 Laudenslager, M. L., Bettinger, T. & Sackett, G. P. (2006). Saliva as a medium for
- assessing cortisol and other compounds in nonhuman primates: collection, assay,
- and examples. In G. P. Sackett, G. C. Ruppenthal, & K. Elias (Eds.). Nursery rearing
- of nonhuman primates in the 21st century (pp. 403–427). Chicago: Springer.
- 883 https://doi.org/10.1007/978-0-387-25640-5_20
- Laudenslager, M. L., Jorgensen, M. J. & Fairbanks, L. A. (2012). Developmental patterns
- of hair cortisol in male and female nonhuman primates: lower hair cortisol levels
- in vervet males emerge at puberty. *Psychoneuroendocrinology*, 37, 1736–1739.
- 887 https://doi.org/10.1016/j.psyneuen.2012.03.015

888	Leung, T. N., Chung, T. K. H., Madsen, G., Lam, P. K., Sahota, D. & Smith, R. (2001). Rate
889	of rise in maternal plasma corticotrophin-releasing hormone and its relation to
890	gestational length. British Journal of Obstetrics and Gynaecology, 108, 527–532.
891	https://doi.org/10.1111/j.1471-0528.2001.00112.x
892	López, J., Wormell, D. & Rodríguez, A. (2001). Preliminary evaluation of the efficacy
893	and safety of a UVB lamp used to prevent metabolic bone disease in pied
894	tamarins Saguinus bicolor at Jersey Zoo. Dodo, 37, 41–49.
895	Macbeth, B. J., Cattet, M. R., Obbard, M. E., Middel, K. & Janz, D. M. (2012). Evaluation
896	of hair cortisol concentration as a biomarker of long-term stress in free-ranging
897	polar bears. Wildlife Society Bulletin, 36, 747–758.
898	https://doi.org/10.1002/wsb.219
899	Mallapur, A. & Choudhury, B. C. (2003). Behavioral abnormalities in captive nonhuman
900	primates. Journal of Applied Animal Welfare Science, 6, 275–284.
901	https://doi.org/10.1207/s15327604jaws0604_2
902	Martin, J. G. A. & Reale, D. (2008). Animal temperament and human disturbance:
903	Implications for the response of wildlife to tourism. Behavioural Processes, 77,

- 904 66–72. https://doi.org/10.1016/j.beproc.2007.06.004
- 905 Martin, L. B. (2009). Stress and immunity in wild vertebrates: timing is everything.
- 906 *General and Comparative Endocrinology*, 163, 70–76.
- 907 https://doi.org/10.1016/j.ygcen.2009.03.008

908	Mason, G. (2010). Species differences in responses to captivity: stress, welfare and the
909	comparative method. Trends in Ecology and Evolution, 25, 713–721.
910	https://doi.org/10.1016/j.tree.2010.08.011
911	Mason, G., Clubb, R., Latham, N. & Vickery, S. (2007). Why and how should we use
912	environmental enrichment to tackle stereotypic behaviour? Applied Animal
913	Behaviour Science, 102, 163–188.
914	https://doi.org/10.1016/j.applanim.2006.05.041
915	McCallister, J. (2005). Behavioural and physiological differences between callitrichid
916	primates. The Queen's University of Belfast, UK (unpublished doctoral thesis).
917	Millspaugh, J. & Washburn, B. (2004). Use of fecal glucocorticoid metabolite measures
918	in conservation biology research: considerations for application and
919	interpretation. General and Comparative Endocrinology 138, 189–199.
920	https://doi.org/10.1016/j.ygcen.2004.07.002
921	Mittermeier, R. A., Boubli, JP., Subirá, R. & Rylands, A. B. (2008). Saguinus bicolor. The
922	IUCN Red List of Threatened Species 2008: e.T40644A10348136.
923	http://dx.doi.org/10.2305/IUCN.UK.2008.RLTS.T40644A10348136.en.
924	Downloaded on 10 August 2017.
925	Morgan, K. & Tromborg, C. (2007). Sources of stress in captivity. Applied Animal

- 926 *Behaviour Science,* 102, 262–302.
- 927 https://doi.org/10.1016/j.applanim.2006.05.032

928	Möstl, E. & Palme, R. (2002). Hormones as indicators of stress. Domestic Animal
929	Endocrinology, 23, 67–74. https://doi.org/10.1016/s0739-7240(02)00146-7
930	Munro, C. & Stabenfeldt, G. (1985). Development of a cortisol enzyme-immunoassay in
931	plasma. <i>Clinical Chemistry</i> , 31, 956–956.
932	Munson, L., Terio, K. A., Worley, M., Jago, M., Bagot-Smith, A. & Marker, L. (2005).
933	Extrinsic factors significantly affect patterns of disease in free-ranging and
934	captive cheetah (Acinonyx jubatus) populations. Journal of Wildlife Diseases, 41,
935	542–548. https://doi.org/10.7589/0090-3558-41.3.542
936	Niebruegge, K. & Porton, I. (2006). The effect of hand rearing on the sexual and
937	maternal competence of three species of lemurs, Varecia variegata, Varecia
938	rubra and Eulemur macaco. In G. P. Sackett, G. C. Ruppenthal and K. Elias (Eds.).
939	Nursery rearing of nonhuman primates in the 21st century (pp. 101–110).
940	Boston, MA: Springer. https://doi.org/10.1007/978-0-387-25640-5_6
941	Pauling, C. D., Lankford, S. E. & Jackson, V. L. (2017). Fecal cortisol levels in scimitar-
942	horned oryx, Oryx dammah, reveal differences between captive environments.
943	Journal of Zoo and Wildlife Medicine, 48, 1049–1057.
944	https://doi.org/10.1638/2016-0087r2.1
945	Pearson, B. L., Reeder, D. M. & Judge, P. G. (2015). Crowding increases salivary cortisol
946	but not self-directed behavior in captive baboons. American Journal of
947	Primatology, 77, 462–467. https://doi.org/10.1002/ajp.22363

948	Pereyra, M. E. & Wingfield, J. C. (2003). Changes in plasma corticosterone and
949	adrenocortical response to stress during the breeding cycle in high altitude
950	flycatchers. General and Comparative Endocrinology, 130, 222–231.
951	https://doi.org/10.1016/s0016-6480(02)00568-3
952	Pinheiro, J., Bates, D., DebRoy, S., Sarkar, D., & R Core Team (2018). nlme: Linear and
953	nonlinear mixed effects models. R package version 3.1-137, https://CRAN.R-
954	project.org/package=nlme
955	Pirovino, M., Heistermann, M., Zimmerman, N., Zingg, R., Clauss, M., Codron, D., Kaup,
956	FJ. & Steinmetz H. W. (2011). Fecal glucocorticoid measurements and their
957	relation to rearing, behavior, and environmental factors in the population of
958	pileated gibbons (Hylobates pileatus) held in European zoos. International
959	Journal of Primatology, 32, 1161–1178. https://doi.org/10.1007/s10764-011-
960	9532-9
961	Pizzutto, C. S., Sgai, M. G. F. G., Francischini, C. P., Viau, P., de Oliveira, C. A. &
962	Guimarães, M. D. B. V. (2015). Physiological validation of enzyme immunoassay
963	of fecal glucocorticoid metabolite levels and diurnal variation measured in
964	captive black-tufted marmoset Callithrix penicillata (Mammalia: Primates:
965	Callitrichidae). Journal of Threatened Taxa, 7, 7234–7242.
966	https://doi.org/10.11609/jott.o4099.7234-42
967	Porton, I. & Niebruegge, K. (2006). The changing role of hand rearing in zoo-based
968	primate breeding programs. In G.P. Sackett, G.C. Ruppenthal and K. Elias (Eds.).

969	Nursery rearing of nonhuman primates in the 21st century (pp. 21–31). Boston,
970	MA Springer. https://doi.org/10.1007/978-0-387-25640-5_2
971	Powell, D. M., Carlstead, K., Tarou, L. R., Brown, J. L. & Monfort, S. L. (2006). Effects of
972	construction noise on behavior and cortisol levels in a pair of captive giant
973	pandas (Ailuropoda melanoleuca). Zoo Biology, 25, 391–408.
974	https://doi.org/10.1002/zoo.20098
975	Powell, D. M. & Watters, J. V. (2017). The evolution of the animal welfare movement in
975 976	Powell, D. M. & Watters, J. V. (2017). The evolution of the animal welfare movement in U.S. zoos and aquariums. <i>Der Zoologische Garten</i> , 86, 219–234.
976	U.S. zoos and aquariums. <i>Der Zoologische Garten</i> , 86, 219–234.
976 977	U.S. zoos and aquariums. <i>Der Zoologische Garten</i> , 86, 219–234. https://doi.org/10.1016/j.zoolgart.2017.04.007

- 981 Price, E. C., & McGrew, W. C. (1991). Departures from monogamy in colonies of captive
- 982 cotton-top tamarins. *Folia Primatologica*, 57, 16–27.
- 983 https://doi.org/10.1159/000156559
- 984 Price, E. C., Payne, C. & Wormell, D. (2016). Why do captive pied tamarins give birth
- 985 during the day? *Zoo Biology*, 35, 487–494. https://doi.org/10.1002/zoo.21325
- 986 Pride, E. (2005). Optimal group size and seasonal stress in ring-tailed lemurs (*Lemur*
- 987 *catta*). Behavioral Ecology, 16, 550–560. https://doi.org/10.1093/beheco/ari025

988 Puterman, E., O'Donovan, A., Adler, N.E., Tomiyama, A.J., Kemeny, M., Wolkowitz,

- 989 O.M. & Epel, E. (2011). Physical activity moderates stressor-induced rumination
- 990 on cortisol reactivity. *Psychosomatic Medicine*, 73, 604–611.
- 991 https://doi.org/10.1097/psy.0b013e318229e1e0
- 992 Quadros, S., Goulart, V. D., Passos, L., Vecci, M. A. & Young, R. J. (2014). Zoo visitor
- 993 effect on mammal behaviour: Does noise matter? *Applied Animal Behaviour*

994 *Science*, 156, 78–84. https://doi.org/10.1016/j.applanim.2014.04.002

995 Queyras, A. & Carosi, M. (2004). Non-invasive techniques for analysing hormonal

996 indicators of stress. *Annali dell'Istituto superiore di sanita*, 40, 211–221.

997 R Core Team (2018). R: A language and environment for statistical computing. Vienna:

998 R Foundation for Statistical Computing. http://www.R-project.org/

- 999 Rajagopal, T., Archunan, G. & Sekar, M. (2011). Impact of zoo visitors on the fecal
- 1000 cortisol levels and behavior of an endangered species: Indian blackbuck
- 1001 (Antelope cervicapra L.). Journal of Applied Animal Welfare Science, 14, 18–32.
- 1002 https://doi.org/10.1080/10888705.2011.527598
- 1003 Ryan, S., Thompson, S. D., Roth, A. M. & Gold, K. C. (2002). Effects of hand-rearing on
- 1004 the reproductive success of western lowland gorillas in North America. *Zoo*
- 1005 *Biology*, 21, 389–401. https://doi.org/10.1002/zoo.10045
- 1006 Saltzman, W., Schultz-Darken, N. J., Wegner, F. H., Wittwer, D. J. & Abbott, D. H.

1007 (1998). Suppression of cortisol levels in subordinate female marmosets:

- 1008 Reproductive and social contributions. *Hormones and Behavior*, 33, 58–74.
- 1009 https://doi.org/10.1006/hbeh.1998.1436
- 1010 Saltzman, W., Liedl, K. J., Salper, O. J., Pick, R. R. & Abbott, D. H. (2008). Post-
- 1011 conception reproductive competition in cooperatively breeding common

1012 marmosets. *Hormones and Behavior*, 53, 274–286.

- 1013 https://doi.org/10.1016/j.yhbeh.2007.10.005
- 1014 Saltzman, W., Prudom, S. L., Schultz-Darken, N. J., Wittwer, D. J. & Abbott, D. H. (2004).
- 1015 Social suppression of cortisol in female marmoset monkeys: role of circulating
- 1016 ACTH levels and glucocorticoid negative feedback. *Psychoneuroendocrinology*,

1017 29, 141–161. https://doi.org/10.1016/s0306-4530(02)00159-2

- 1018 Savage, A., Giraldo, L. H., Soto, L. H. & Snowdon, C. T. (1996). Demography, group
- 1019 composition, and dispersal in wild cotton-top tamarin (*Saguinus oedipus*) groups.
- 1020 American Journal of Primatology, 38, 85–100. https://doi.org/10.1002/(sici)1098-
- 1021 2345(1996)38:1%3C85::aid-ajp7%3E3.0.co;2-p
- 1022 Schaffner, C. M. & Caine, N. G. (2000). The peacefulness of cooperatively breeding
- 1023 primates. In F. Aureli & F. de Waal (Eds.), Natural conflict resolution (pp. 155–
- 1024 172). Oakland: University of California Press.
- 1025 Schoof, V. A., Jack, K. M. & Ziegler, T. E. (2014). Male response to female ovulation in
- 1026 white-faced capuchins (*Cebus capucinus*): variation in fecal testosterone,
- 1027 dihydrotestosterone, and glucocorticoids. International Journal of
- 1028 Primatology, 35, 643–660. https://doi.org/10.1007/s10764-013-9742-4

1029	Schumann, K., Guenther, A., Jewgenow, K. & Trillmich, F. (2014). Animal housing and
1030	welfare: effects of housing conditions on body weight and cortisol in a medium-
1031	sized rodent (Cavia aperea). Journal of Applied Animal Welfare Science, 17, 111–
1032	124. https://doi.org/10.1080/10888705.2014.884407
1033	Shepherdson, D. J., Carlstead, K. C. & Wielebnowski, N. (2004). Cross-institutional
1034	assessment of stress responses in zoo animals using longitudinal monitoring of
1035	faecal corticoids and behaviour. Animal Welfare, 13, S105–S114.
1036	Shepherdson, D., Lewis, K. D., Carlstead, K., Bauman, J. & Perrin, N. (2013). Individual
1037	and environmental factors associated with stereotypic behavior and fecal
1038	glucocorticoid metabolite levels in zoo housed polar bears. Applied Animal
1039	Behaviour Science, 147, 268–277.
1040	https://doi.org/10.1016/j.applanim.2013.01.001
1041	Sherwen, S. L., Harvey, T. J., Magrath, M. J., Butler, K. L., Fanson, K. V. & Hemsworth, P.
1042	H. (2015). Effects of visual contact with zoo visitors on black-capped capuchin
1043	welfare. Applied Animal Behaviour Science, 167, 65–73.
1044	https://doi.org/10.1016/j.applanim.2015.03.004
1045	Simpkins, K., Routh, A., Wormell, D. & Price, E. (2013). The effects of noise on faecal
1046	cortisol and behaviour in off-show pied tamarins (Saguinus bicolor) at Durrell
1047	Wildlife Conservation Trust. Royal Veterinary College, London (unpublished MSc

1048 dissertation).

1049	Smith, T. E. & French, J. A. (1997a). Psychosocial stress and urinary cortisol excretion in
1050	marmoset monkeys. Physiology and Behavior, 62, 225–232.
1051	https://doi.org/10.1016/s0031-9384(97)00103-0
1052	Smith, T. E. & French, J. A. (1997b). Social and reproductive conditions modulate
1053	urinary cortisol excretion in black tufted-ear marmosets (Callithrix kuhli).
1054	American Journal of Primatology, 42, 253–267.
1055	https://doi.org/10.1002/(sici)1098-2345(1997)42:4%3C253::aid-
1056	ajp1%3E3.0.co;2-w
1057	Smith, T. E., McCusker, C. M., Stevens, J. M. & Elwood, R. W. (2015). Patterns of
1058	behaviour, group structure and reproductive status predict levels of
1059	glucocorticoid metabolites in zoo-housed ring-tailed lemurs, Lemur catta. Folia
1060	<i>Primatologica</i> , 86, 506–524. https://doi.org/10.1159/000442587
1061	Smith, T.E., McGreer-Whitworth, B. and French, J.A. (1998). Close proximity of the
1062	heterosexual partner reduces the physiological and behavioral consequences of
1063	novel-cage housing in black tufted-ear marmosets (Callithrix kuhli). Hormones
1064	and Behavior, 34, 211–222. https://doi.org/10.1006/hbeh.1998.1469
1065	Smithyman, J. (2012). Wasting marmoset syndrome in pied tamarins (Saguinus
1066	bicolor). Durrell Wildlife Conservation Trust, Jersey (Unpublished report).
1067	Sousa, M. B. C. & Ziegler, T. E. (1998). Diurnal variation on the excretion patterns of
1068	fecal steroids in common marmoset (Callithrix jacchus) females. American

1069	Journal of Primatology, 46, 105–117. https://doi.org/10.1002/(sici)1098-
1070	2345(1998)46:2%3C105::aid-ajp1%3E3.3.co;2-j
1071	Species360 (2018). Species360 Zoological Information Management System (ZIMS)
1072	(2018), zims.Species360.org.
1073	Starling, A.P., Charpentier, M.J., Fitzpatrick, C., Scordato, E.S. & Drea, C.M. (2010).
1074	Seasonality, sociality, and reproduction: long-term stressors of ring-tailed lemurs
1075	(Lemur catta). Hormones and Behavior, 57, 76–85.
1076	https://doi.org/10.1016/j.yhbeh.2009.09.016
1077	Steinbrecher, F. (2016). Factors influencing the vigilance, vocalization and scent
1078	marking behaviour of captive pairs of pied tamarins (Saguinus bicolor). Anglia
1079	Ruskin University (unpublished MSc dissertation).
1080	Strier, K.B., Lynch, J.W. and Ziegler, T.E. (2003). Hormonal changes during the mating
1081	and conception seasons of wild northern muriquis (Brachyteles arachnoides
1082	hypoxanthus). American Journal of Primatology, 61, 85–99.
1083	https://doi.org/10.1002/ajp.10109
1084	Surbeck , M., Deschner, T., Weltring, A. & Hohmann, G. (2012) Social correlates of
1085	variation in urinary cortisol in wild male bonobos Pan paniscus. Hormones and

1086 Behavior, 62, 27–35. https://doi.org/10.1016/j.yhbeh.2012.04.013

1087	Tamashiro, K., Nguyen, M. & Sakai, R. (2005). Social stress: From rodents to primates.
1088	Frontiers in Neuroendocrinology, 26, 27–40.

1089 https://doi.org/10.1016/j.yfrne.2005.03.001

- 1090 Tardif, S. D., Carson, R. L., & Gangaware, B. L. (1986). Comparison of infant care in
- 1091 family groups of the common marmoset (*Callithrix jacchus*) and the cotton-top
- 1092 tamarin (*Saguinus oedipus*). *American Journal of Primatology*, 11, 103–110.
- 1093 https://doi.org/10.1002/ajp.1350110202
- 1094 Tardif, S. D., Richter, C. B. & Carson, R. L. (1984). Effects of sibling-rearing experience

1095 on future reproductive success in two species of Callitrichidae. *American Journal* 1096 of *Primatology*, 6, 377–380. https://doi.org/10.1002/ajp.1350060408

1097 Teixeira, R. R., Díaz, M. M., da Silva Santos, T. V., Bernardes, J. T. M., Peixoto, L. G.,

1098 Bocanegra, O. L., Neto, M. B. & Espindola, F. S. (2015). Chronic stress induces a

1099 hyporeactivity of the autonomic nervous system in response to acute mental

1100 stressor and impairs cognitive performance in business executives. *PloS One*,

1101 10(3), e0119025. https://doi.org/10.1371/journal.pone.0119025

1102 Thomas, R., Lello, J., Medeiros, R., Pollard, A., Robinson, P., Seward, A., Smith, J.,

1103 Vafidis, J., & Vaughan, I. (2017). Data analysis with R statistical software. Cardiff:
1104 Eco-explore.

Traylor-Holzer, K., Leus, K. & Byers, O. (2018). Integrating ex situ management options
as part of a one plan approach to species conservation. In B. A. Minteer, J.
Maienschein, J. P. Collins, & G. Rabb (Eds.). The ark and beyond: The evolution of

1108	zoo and aquarium conservation (pp. 129–141). Chicago: University of Chicago
1109	Press. https://doi.org/10.7208/chicago/9780226538631.003.0011
1110	Trevisan, C., Montillo, M., Prandi, A., Mkupasi, E. M., Ngowi, H. A. & Johansen, M. V.
1111	(2017). Hair cortisol and dehydroepiandrosterone concentrations in naturally
1112	Taenia solium infected pigs in Tanzania. General and Comparative Endocrinology,
1113	246, 23–28. https://doi.org/10.1016/j.ygcen.2017.03.007
1114	Vermeer, J. & Devreese, L. (2015). Birth sex ratio, infant mortality and rearing type in
1115	captive western lowland gorillas. Journal of Zoo and Aquarium Research, 3, 6–10.
1116	Wark, J. D., Amendolagine, L., Lukas, K. E., Kuhar, C. W., Dennis, P. M., Snowdon, C. T.,
1117	Schoffner, T. & Schook, M. W. (2016). Fecal glucocorticoid metabolite responses
1118	to management stressors and social change in four species of callitrichine
1119	monkeys. <i>Primates</i> , 57, 267–277. https://doi.org/10.1007/s10329-016-0514-6
1120	Wasser, S. K., Thomas, R., Nair, P. P., Guidry, C., Southers, J., Lucas, J., Wildt, D. E., &
1121	Monfort, S. L. (1993). Effects of dietary fibre on faecal steroid measurements in
1122	baboons (Papio cynocephalus cynocephalus). Reproduction, 97, 567–574.
1123	https://doi.org/10.1530/jrf.0.0970569
1124	Wormell, D. (2000). Management guidelines for pied tamarins Saguinus bicolor bicolor.
1125	<i>Dodo,</i> 36, 87.
1126	Wormell D., & Brayshaw, M. (2000). The design and redevelopment of New World

1127 primate accommodation at Jersey Zoo: a naturalistic approach. *Dodo*, 36, 9–19.

1128	Wormell, D., Brayshaw, M., Price, E. & Herron, S. (1996). Pied tamarins Saguinus
1129	bicolor at the Jersey Wildlife Preservation Trust: management, behaviour and
1130	reproduction. <i>Dodo, Journal of the Wildlife Preservation Trusts</i> , 32, 76–97.
1131	Ziegler, T. E. (2013). Social effects via olfactory sensory stimuli on reproductive
1132	function and dysfunction in cooperative breeding marmosets and tamarins.
1133	American Journal of Primatology, 75, 202–211.
1134	https://doi.org/10.1002/ajp.22061
1135	Ziegler, T. E., Scheffler, G. & Snowdon, C. T. (1995). The relationship of cortisol levels
1136	to social environment and reproductive functioning in female cotton-top

1137 tamarins, *Saguinus oedipus. Hormones and Behavior*, 29, 407–424.

1138 https://doi.org/10.1006/hbeh.1995.1028

	Saliva sam	ples	Fecal sam	Fecal samples	
	No. of	No. of	No. of	No. of	
	samples	individuals	samples	individuals	
Sex					
Female	78	16	102	12	
Male	136	20	125	15	
Rearing history					
Hand reared	36	5	48	5	
Parent reared	178	31	179	22	
Group composition*					
Family	68	17	140	18	
Pair	37	10	42	6	
Single sex	21	2	19	2	
Mixed species	87	15	26	4	
Social status*					
Dominant	147	27	151	17	
Subordinate	67	11	76	10	
Enclosure type*					
Off-show	159	28	134	21	
On-show	55	9	93	12	
Pregnancy state (breeding females only)					
Early pregnancy	1	1	14	3	
Late pregnancy	3	2	3	2	
Not pregnant	26	6	47	7	
Total	214	36	227	27	

Table 1. Sample sizes for each factor included in the GLMMs.

*Number of individuals may sum to more than overall total as some tamarins contributed samples under more than one condition.

Model	df	AICc	ΔAIC_{c}	Weight
Enclosure + rearing + sex + social status + weight +	9	710.39	0.00	0.14
rearing:sex				
Enclosure + social status + weight	6	710.74	0.36	0.11
Enclosure + rearing + sex + weight + rearing:sex	8	710.77	0.38	0.11
Enclosure + rearing + weight	6	710.91	0.53	0.10
Enclosure + weight	5	711.12	0.73	0.09
Enclosure + rearing + social status + weight	7	711.27	0.88	0.09
Enclosure + rearing + sex + social status + age + weight +	10	711.61	1.22	0.07
rearing:sex				
Enclosure + rearing + sex + weight	7	711.82	1.43	0.07
Enclosure + sex	5	712.05	1.66	0.06
Enclosure + sex + social status + weight	7	712.27	1.89	0.05
Enclosure + rearing + sex + social status + weight	8	712.31	1.93	0.05
	1			

9

712.33

1.94

0.05

Table 2. Models with $\Delta AIC_c < 2$ for log cortisol in fecal samples.

Enclosure + social status + group composition + weight

Table 3. Estimates, standard errors and 95% confidence intervals for predictor

variables in averaged model for log fecal cortisol. The first level listed for each variable

is the reference level.

			Confidence intervals		
Variable	Estimate (β)	SE	2.5 %	97.5 %	
Enclosure (offshow–onshow)	-0.5197	0.2275	-0.9680	-0.0715	
Weight	-0.5482	0.2693	-1.1132	-0.0520	
Rearing (hand-parent)	-0.4181	0.3497	-1.1071	0.2709	
Social status (dominant-subordinate)	-0.4585	0.2925	-1.0348	0.1179	
Age	-0.3104	0.3130	-0.9274	0.3067	
Sex (female–male)	0.2794	0.2587	-0.2305	0.7893	
Rearing:sex	1.3365	0.6609	0.0339	2.6392	
Group composition (family-mixed)	0.7580	0.3695	0.0297	1.4864	
Group composition (family-pair)	0.3760	0.3221	-0.2591	1.0120	
Group composition (family-single sex)	-0.1267	0.4640	-1.0414	0.7880	
Group composition (mixed-pair)	-0.3821	0.4431	-1.2555	0.4913	
Group composition (mixed-single sex)	-0.8848	0.5433	-1.9557	0.1862	
Group composition (pair-single sex)	-0.5027	0.5116	-1.5111	0.5058	

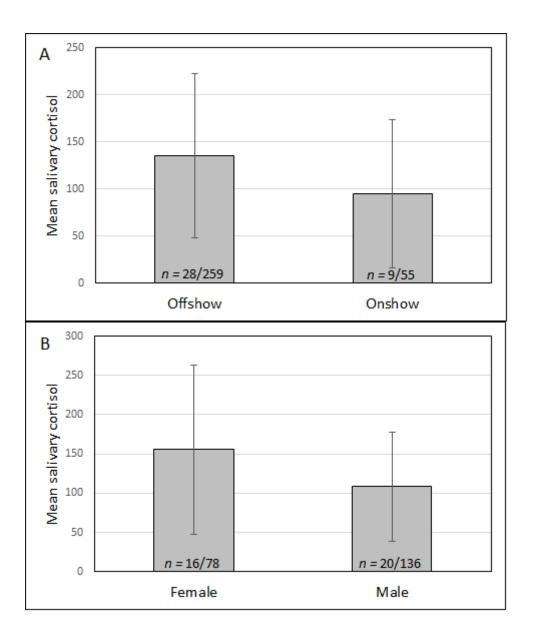


Figure 1. A: Mean salivary cortisol in pied tamarins housed in different enclosure types. B: Mean salivary cortisol in male and female pied tamarins. Other factors did not appear in the final GLMM and are not illustrated. Means were back-transformed from mean log cortisol values. Vertical bars represent coefficient of variation. Sample sizes (number of individuals/number of samples) are given at the base of each bar.

Price - 61

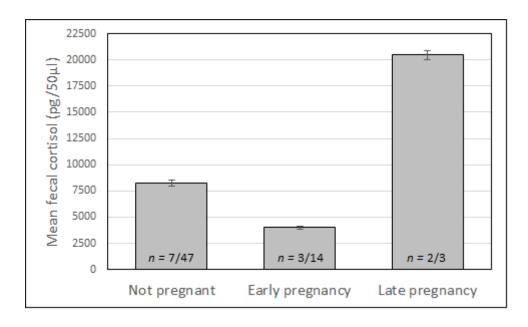


Figure 2. Effect of pregnancy stage on mean fecal cortisol in breeding females. Means were back-transformed from mean log cortisol values. Vertical bars represent coefficient of variation. Sample sizes (number of individuals/number of samples) are given at the base of each bar.

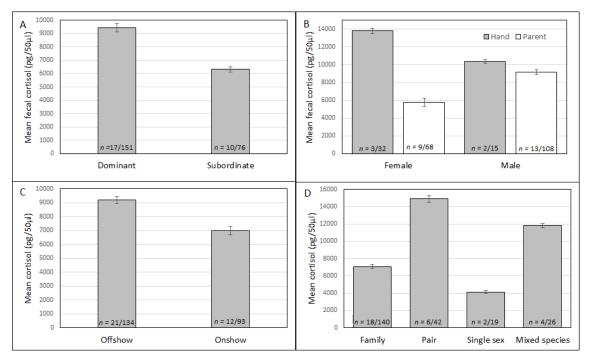


Figure 3. A: Mean fecal cortisol in dominant and subordinate pied tamarins. B: Effect of sex and rearing history on mean fecal cortisol in pied tamarins. C: Mean fecal cortisol in pied tamarins housed in different enclosure types. D: Mean fecal cortisol in pied tamarins housed in different group types. Means were back-transformed from mean log cortisol values; vertical bars represent coefficient of variation. Sample sizes (number of individuals/number of samples) are given at the base of each bar.

Price - 62

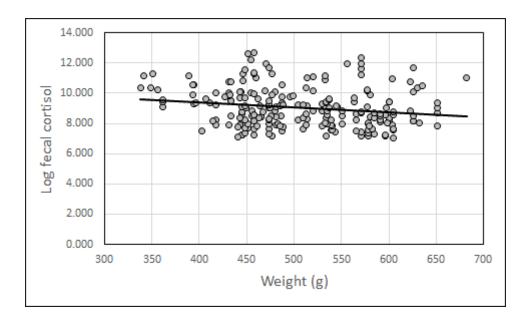


Figure 4. Relationship between log-transformed fecal cortisol levels and weight in pied tamarins.



Figure 5. Large on-show enclosure for pied tamarins at Jersey Zoo, showing stand-off barrier and planting to reduce contact with visitors. A pied tamarin is visible in the center of the image.