

# UCLA

## UCLA Previously Published Works

### Title

Are demographic correlates of white-faced capuchin monkey (*Cebus capucinus*) "Gargle and Twargle" vocalization rates consistent with the infanticide risk assessment hypothesis?

### Permalink

<https://escholarship.org/uc/item/8781s65x>

### Journal

American Journal of Primatology, 84(1)

### ISSN

0275-2565

### Authors

Duchesneau, Alexa  
Edelberg, Daniel G  
Perry, Susan E

### Publication Date

2022

### DOI

10.1002/ajp.23344

Peer reviewed

1

2

1 Title: Are demographic correlates of White-faced Capuchin Monkeys

2 (*Cebus capucinus*) “Gargle and Twargle” Vocalization Rates consistent

3 with the Infanticide Risk Assessment Hypothesis?

4

5 Running title: Capuchins test bonds via vocalizations

6

7

8 Authors: Alexa Duchesneau<sup>1</sup>, Daniel G. Edelberg<sup>2</sup>, Susan E. Perry<sup>3,4</sup>

9 <sup>1</sup>Department of Anthropology, Yale University, <sup>2</sup>Department of

10 Mathematics, Yale University, <sup>3</sup>Department of Anthropology, University

11 of California at Los Angeles, <sup>4</sup>Behavior, Evolution and Culture Program,

12 University of California at Los Angeles

13

14 Corresponding author: Susan Perry, Dept. of Anthropology, UCLA, 341

15 Haines Hall, 375 Portola Plaza, Los Angeles, CA 90095.

16 [sperry@anthro.ucla.edu](mailto:sperry@anthro.ucla.edu)

17 ORCID ID: [0000-0001-5306-5383](https://orcid.org/0000-0001-5306-5383)

18

19 Submitted to the American Journal of Primatology, subsequently published at

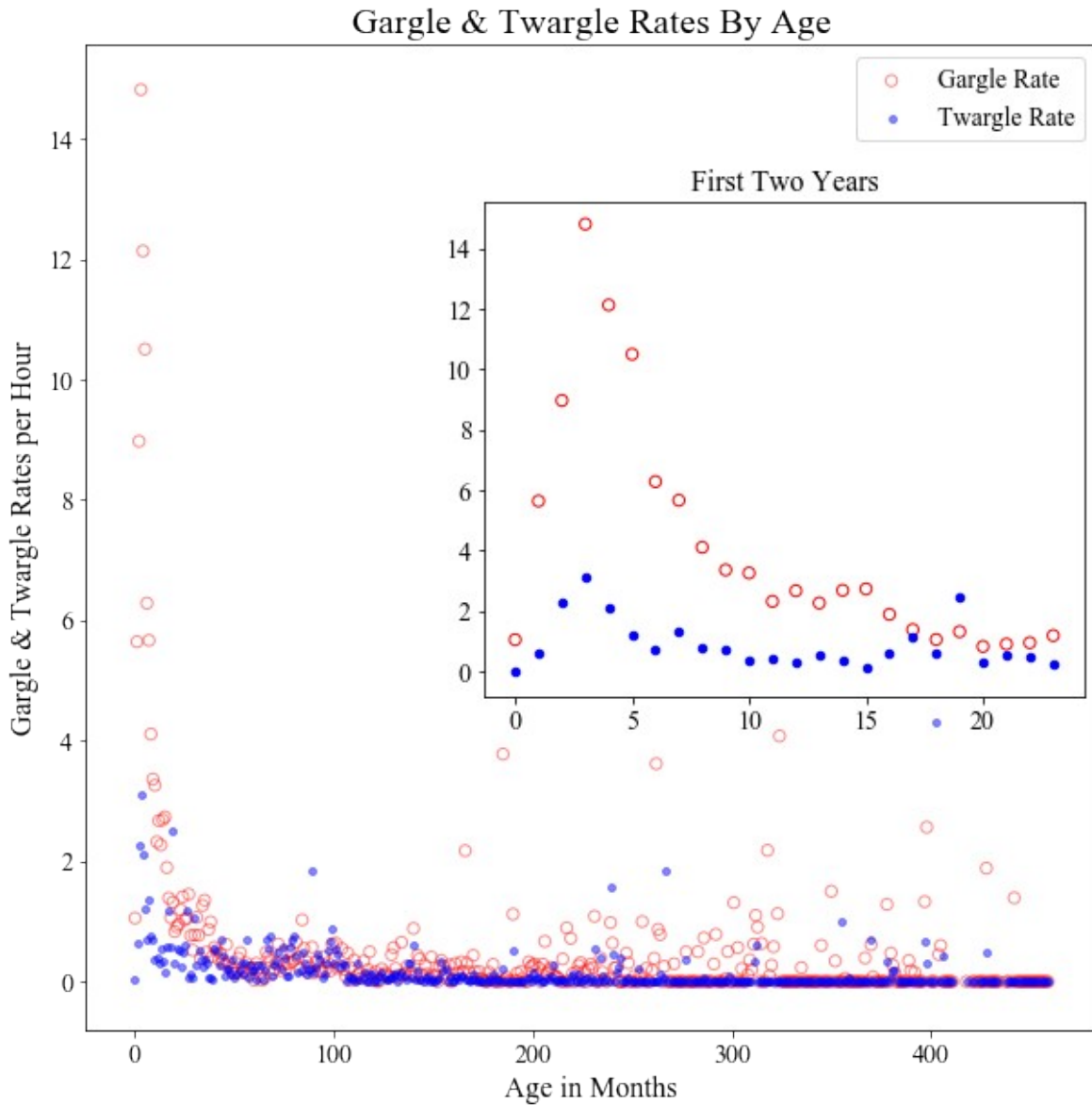
20 DOI:10.1002/ajp.23344

3

4

21 **Graphical Abstract**

22 The scatterplot displays average gargle (open red circles) and twargle (closed blue circles) rate data by age in months. The subplot displays average gargle and  
 23 gargle and  
 24 twargle rates for individuals 24 months of age and younger. Individuals gargle and  
 25 twargle the most during infancy, when risk of infanticide is the highest.



26

5

6

27 **Abstract:**

28 Zahavi's "Bond Testing Hypothesis" (1977a) states that irritating  
29 stimuli are used to elicit honest information from social partners  
30 regarding their attitudes towards the relationship. Two elements of the  
31 *C. capucinus* vocal repertoire, the "gargle" and "twargle," have been  
32 hypothesized to serve such a bond-testing function (Perry & Manson,  
33 2008). The greatest threat to *C. capucinus* infant survival, and to adult  
34 female reproductive success, is infanticide perpetrated by alpha males  
35 (Perry, 2012). Thus, we predicted that infants (<8 months), pregnant  
36 females and females with infants would gargle/twargle at higher rates  
37 than the rest of the population, directing these vocalizations primarily to  
38 the alpha male. Over 16 years, researchers collected data via focal  
39 follows in 11 habituated groups of wild capuchins in Lomas Barbudal,  
40 Costa Rica. Our hypothesis was mainly supported. Infants and females  
41 with infants (<8 months) vocalized at higher rates than the rest of the  
42 population. Pregnant females did not vocalize at high rates. Infants (age  
43 8-23 mo.) were the only target group that vocalized more when the  
44 alpha male was not their father. Monkeys gargled and twargled most  
45 frequently towards the alpha male, who is both the perpetrator of  
46 infanticide and the most effective protector against potentially  
47 infanticidal males.

48

7

8

49 **Key Words:** *Cebus capucinus*, *Zahavian bond tests*, *vocal*

50 *communication*, *infanticide*

51

9

10

52 **Introduction**

53 Much of the nonhuman primate vocalization literature focuses on how  
54 vocalizations broadcast, rather than elicit, information (Bergman,  
55 Beehner, Painter, & Gustison, 2019; Elowson, Snowdon, & Lazaro-Perea,  
56 1998; Gros-Louis, 2002, 2006; Gros-Louis et al., 2008; Locke, 2001;  
57 Maynard Smith, 1982; Schamberg, Cheney, Clay, Hohmann, & Seyfarth,  
58 2017; Seyfarth & Cheney, 1986). Researchers may overlook calls  
59 functioning as “Zahavian bond tests” when centering research on vocal  
60 broadcasting rather than elicitation (Zahavi 1977a). A Zahavian Bond  
61 Test is the engagement in a risky behavior, or stimulus, imposed by a  
62 “tester” to elicit an honest response regarding the sentiments of the  
63 recipient towards the tester (Zahavi, 1977b; Zahavi & Zahavi, 1977). In  
64 this paper, we examine the possibility that white-faced capuchins (*Cebus*  
65 *capucinus*) use two calls from their vocal repertoire, the “gargle” and  
66 “twargle,” to elicit information about a recipient’s sentiments towards  
67 the caller, especially if the caller faces infanticide risk.

68 Research suggests that white-faced capuchins navigate  
69 relationships by testing and reinforcing bonds via non-vocal signals  
70 (Manson, 1999; Perry, 2011; Perry et al., 2003; Perry & Manson, 2008). A  
71 bond, defined by Zahavi (1977a) as a “special relationship between two  
72 individuals,” may form between parents-offspring, sexual partners or  
73 group members and may change through time (Zahavi, 1977a).

11

12

74 Zahavian bond tests promote honest communication within a dyad, as  
75 the tester aims to discern the attitudes of the recipient towards the  
76 tester (Taylor, 2014; Zahavi, 1977a). A neutral or positive response from  
77 the recipient (e.g., producing an affiliative vocalization, gesturing a  
78 greeting, continuing with the same behavior, etc.) may indicate interest  
79 in relationship investment, and a negative response or premature  
80 termination of the stimuli (e.g., hitting, walking away, biting, etc.) may  
81 indicate disinterest in relationship investment (Smuts & Watanabe,  
82 1990; Zahavi, 1977a). For example, old adult male olive baboons (*Papio*  
83 *anubis*), as compared to young adult males, generally complete bond  
84 testing behaviors, indicating relationship investment. The Zahavian Bond  
85 Testing Hypothesis would predict this outcome, as old adult males are  
86 reliant on allies for coalitionary support, whereas young adult males are  
87 not (Smuts & Watanabe, 1990).

88         A signal is more likely to be trusted if it is costly or risky to  
89 produce, because it would be unprofitable for unmotivated individuals to  
90 produce such signals (Zahavi, 1977a; Zahavi & Zahavi, 1977). Signaling  
91 does not result in cooperation when individuals do not trust information  
92 conveyed by their partner (Silk, Kaldor, & Boyd, 2000). Some group-  
93 living primates, such as chacma baboons (*Papio ursinus*) and rhesus  
94 macaques (*Macaca mulatta*) produce honest signals and responses to  
95 coordinate coalitionary behavior (Silk et al., 2000; Silk, Seyfarth, &

13

14

96 Cheney, 2016). Dishonest signalers potentially incur punishment from  
97 group members, resulting in negative long-term consequences for  
98 perpetrators (Cheney & Seyfarth, 2018; Poole, 1989; Silk et al., 2000,  
99 2016).

100         White-faced capuchins are known to produce risky signals and  
101 perform Zahavian Bond Tests (Manson, 1999; Perry, 2011; Perry et al.,  
102 2003; Perry & Manson, 2008; Perry & Smolla, 2020). Females test bonds  
103 by holding allies' infants in risky acts of trust (Manson, 1999), and dyads  
104 of all age-sex classes engage in risky rituals. For example, individuals  
105 stick sharp and dirty objects or body parts (such as wood chips, fingers,  
106 or feet) into another individual's eye-socket, nose or mouth (Perry, 2011,  
107 2012; Perry et al., 2003; Perry & Smolla, 2020).

108         We expect white-faced capuchins to test bonds with individuals  
109 with whom they have important relationships, such as parents,  
110 alloparents, allies and adult males. Relationships with alpha males are  
111 arguably some of the most critical relationships in white-faced capuchin  
112 societies, as alpha males are capable of providing great benefits (e.g.,  
113 resources or coalitionary support) or imposing tremendous costs (e.g.,  
114 stress or death) on individuals (Perry, 2012). Alpha males benefit from  
115 relationships with females for reproductive opportunities largely  
116 unavailable to subordinate males (Perry, 1997, 1998, 2012).



15

16

117 Subordinate males also, but to a lesser degree, provide invaluable  
118 resources to group members, such as coalitionary support and  
119 protection against predators and extra-group males (Perry, 1997, 2012).  
120 However, they can also pose a threat to capuchin societies by  
121 overthrowing alpha males and subsequently destabilizing social  
122 interactions, altering group behavior and committing infanticide (Jack &  
123 Fedigan, 2008; Perry, 2012; Perry et al., 2003; Perry, Godoy, & Lammers,  
124 2012; Perry, Godoy, Lammers, & Lin, 2017). Infanticide, i.e. the killing of  
125 unweaned offspring, creates breeding opportunities for perpetrators  
126 (Hrdy, 1979; Palombit, 1999; Perry et al., 2012). Alpha males are the  
127 only known infanticide perpetrators at the Lomas Barbudal site (Perry,  
128 2012), yet infanticide is the leading cause of infant white-faced capuchin  
129 deaths (49%-82% of deaths occurring in the wake of takeover events  
130 and 12%-18% during peaceful periods) and poses the largest threat to  
131 female reproductive success (Fedigan, 2003; Perry, 2012; Perry et al.,  
132 2012, 2017).

133 It is critical for individuals, especially adult females and infants, to  
134 sensibly navigate bonds with the alpha male, given the benefits and  
135 costs of these relationships. Individuals profit from accurately assessing  
136 if the alpha male is either: 1) willing to invest in the bond and provide  
137 coalitionary support, or 2) unwilling to invest in the bond and potentially  
138 threaten an infant's life. We present and test the "Infanticide Risk

17

18

139 Assessment Hypothesis” (Perry & Manson, 2008), suggesting that  
140 individuals use Zahavian Bond Tests to assess infanticide risk. We  
141 hypothesize that white-faced capuchins use two elements of their vocal  
142 repertoire, the “gargle” and the “twargle,” to assess relationship quality.  
143 We predict that individuals will gargle and twargle more frequently when  
144 the recipient of these vocalizations poses infanticide risk to the  
145 vocalizer.

146 Below is a list of predictions (in italics) that can be derived from  
147 the Assessment of Infanticide Risk Hypothesis, followed by clarification  
148 of the assumptions underlying each prediction. See Table 1 for each  
149 prediction’s definitions, data sets and statistical approaches.

150 P.1. *Infants will gargle and twargle at higher rates than the rest of*  
151 *the population.* Unweaned infants should be highly motivated to test  
152 bonds, because they face the highest infanticide risk (Perry et al., 2012).  
153 They can more accurately navigate social situations knowing (a) if the  
154 alpha male will protect them against infanticidal males (in which case  
155 they should maintain proximity to him), (b) if the alpha male poses a  
156 threat to them (in which case they should avoid him) and (c) who may  
157 provide coalitionary support (in which case they should affiliate with  
158 those individuals). Infants will gargle and twargle at high rates until they  
159 are largely weaned, and their mothers can conceive again.

19

20

160           P.2. *Pregnant females will gargle and twargle at higher rates than*  
161 *the rest of the: a) adult female population and b) adult females who are*  
162 *not pregnant and do not have an infant <8 months old. They benefit*  
163 *from knowing (a) if the alpha male will protect their expected offspring*  
164 *(in which case they should maintain proximity to him when the baby is*  
165 *born and present), (b) if the alpha male poses a threat to their expected*  
166 *offspring (in which case they should avoid him when the baby is born*  
167 *and present), and (c) who may provide coalitionary support (in which*  
168 *case they should affiliate with those individuals). Pregnant females will*  
169 *gargle and twargle throughout their pregnancy, especially as the*  
170 *birthing event approaches.*

171           P.3. *Females with infants (<8 months) will gargle and twargle at*  
172 *higher rates than the rest of the: a) adult female population and b) adult*  
173 *females who are not pregnant and do not have an infant <8 months old.*  
174 *They benefit from knowing (a) if the alpha male will protect their*  
175 *offspring (in which case they should maintain proximity to him when the*  
176 *baby is near), (b) if the alpha male poses a threat to their offspring (in*  
177 *which case they should avoid proximity to him when the baby is near),*  
178 *and (c) who may provide coalitionary support (in which case they should*  
179 *affiliate with those individuals). Females are expected to frequently test*  
180 *bonds with the alpha male until the end of the weaning period and*  
181 *especially during the three months following the birthing event when*

21

22

182 lactation has the largest impact on a female's ability to conceive again  
183 (Perry, 2012; Treves, 2000; Van Schaik & Dunbar, 1990).

184       P.4. *Individuals will gargle and twargle more when the current*  
185 *alpha male is not the offspring's father*, because individuals should be  
186 motivated to assess bonds with more group members for coalitionary  
187 support if it is uncertain that the alpha male will provide coalitionary  
188 support or pose infanticide risk (Treves, 2000).

189       P.5. *Individuals will gargle and twargle more to the alpha male*  
190 *than to any other monkey in the group*, because alpha males are the  
191 best source of coalitionary support against future potentially infanticidal  
192 alpha males (Perry, 2012). Additionally, alpha males who have recently  
193 acquired tenure (within one calendar year) (Perry, 1998) are the highest  
194 source of infant mortality and threaten female reproductive success  
195 (Perry, 1997, 2012).

196

23

24

197 Table 1. Target groups, reference groups, model/test and data for each  
 198 prediction.

Prediction	Target Group	Reference Group	Model/Statistical Test	Data Type
P.1.	Infants (<8 months)	Rest of population	NB Model	Gargle and twargle counts and hours
P.2.	Pregnant females	(2a) Non-pregnant adult females (2b) Adult females who are not pregnant and do not have an infant <8 months old	NB Model	Gargle and twargle counts and hours
P.3.	Females with young infants (<8 months)	(3a) Adult females without infants (<8 months) (3b) Adult females who are not pregnant and do not have an infant <8 months old	NB Model	Gargle and twargle counts and hours
P.4.	(4a) Infants (<24 months) whose father is not the alpha male (4b) Pregnant females whose fetus is not sired by the alpha male (4c) Females with infants (<8 months) not sired by the alpha male	(4a) Infants (<24 months) whose father is the alpha male (4b) Pregnant females whose fetus is sired by the alpha male (4c) Females with infants (<8 months) sired by the alpha	NB Model	Gargle and twargle counts and hours

25  
26

		male		
P.5.	All individuals	N/A	Binomial Test	Gargle and twargle count proportions directed towards the alpha male

199 **Methods**

200 **(a) Ethical Note**

201       The protocol and procedures were ethically reviewed and  
 202 approved by UCLA’s Animal Research Committee (ARC), which ensures  
 203 compliance with the US NRC’s Guide for the Care and Use of Laboratory  
 204 Animals, the US PHS’s policy on Humane Care and Use of Laboratory  
 205 Animals, and the Guide for the Care and Use of Laboratory Animals; the  
 206 ARC approved protocols relevant to this project are #1996-122, 2005-  
 207 084, 2016-022 (plus various renewals of these). This work was  
 208 conducted with appropriate permission from the Costa Rican authorities  
 209 (SINAC, MINAET, and CONAGEBIO), which granted permits for data  
 210 collection and procedures. All field work complied with Costa Rican law,  
 211 the ASP’s principles for ethical treatment of non-human primates, and  
 212 the code of best field practices for field primatology.

213 **(b) Study System**

214       Data presented were collected from 2002-2018 on 11 groups of  
 215 wild, well-habituated white-faced capuchin monkeys (*Cebus capucinus*)

27

28

216 at the Lomas Barbudal Biological Reserve and surrounding forest  
217 (latitude: 10.510, longitude: -85.380). White-faced capuchins are large-  
218 brained, long-living New World monkeys (Perry, 2012). They reside in  
219 stable multi-female, multi-male, female-philopatric groups ranging in  
220 size from 7-30 individuals (Perry, 2012). Males generally disperse to  
221 neighboring groups around the time of maturity, alone or in groups of 2-  
222 8 individuals (Jack & Fedigan, 2004, 2008; Perry, 2012). White-faced  
223 capuchins have a wide range of learned and species-typical vocal and  
224 gestural behaviors (Gros-Louis et al., 2008). More information about the  
225 site, population and methods can be found in (Frankie et al., 1988; Perry  
226 et al., 2012).

### 227 **(c) Description of the Behavior**

228       The gargle vocalization is a loud, raspy, guttural, broad-band  
229 vocalization generally produced in bouts in close range (~5-10 m) of the  
230 targeted individual (Gros-Louis et al., 2008; Perry, 1998). Gargles are  
231 produced by individuals in all age-sex classes but rarely by adult males  
232 (Gros-Louis et al., 2008; Perry, 1998). Gargles are one of the first  
233 vocalizations infants produce during the first month of life (Gros-Louis et  
234 al., 2008), suggesting an urgency in an individual's ability to produce the  
235 vocalization. Individuals have been observed gargling to group members  
236 in most age-sex classes (Perry, 1998).

29

30

237           The twargle vocalization begins with high-pitched trill sounds and  
238 cascades into low, raspy gargle sounds (Gros-Louis et al., 2008). Trills  
239 are high-pitched vocalizations generally produced in bouts during  
240 affiliative social situations or travel (Gros-Louis, 2002; Gros-Louis et al.,  
241 2008). Individuals in most age-sex classes twargle (Gros-Louis et al.,  
242 2008; Table 2).

243           Gargles and twargles are primarily produced while resting,  
244 travelling or positively affiliating with the recipient (Gros-Louis et al.,  
245 2008). The recipient varies in their reaction to the vocalizer; they may  
246 leave, ignore, act affiliatively (e.g. by grooming or receiving grooming  
247 from the vocalizer) or act aggressively (e.g. by hitting or pushing the  
248 vocalizer).

249           We performed separate analyses on gargles and twargles,  
250 although they are often produced in conjunction. Gargles and twargles  
251 are acoustically distinct, production rates are distributed differently  
252 across age-sex classes (Gros-Louis et al., 2008), and there is no prior  
253 analysis suggesting that they fulfill the same function.

#### 254 **(d) Data Collection**

255           Data were collected using a strict behavioral focal follow protocol.  
256 Focal follows were primarily 10 minutes long (86% of the time spent  
257 conducting focal follows was during 10-minute follows) conducted across  
258 demographics; however, neither individuals nor age-sex classes were



31

32

259 evenly sampled (Table 2; Table S1). At least one research assistant in  
260 the field was required to accurately identify all individuals within a group  
261 before collecting data. Monthly coding, vocalization and data collection  
262 tests were mandated to ensure efficiency and accuracy at using the  
263 behavioral coding scheme. Data were not included in analyses unless at  
264 least one assistant in the field per day could accurately collect data.  
265 More information on the field site protocols can be found in (Perry et al.,  
266 2012).

267

33

34

268 Table 2. Column 1 refers to target populations delineated by age and  
 269 developmental life history stage. “Gargle Count” and “Twargle Count” refer to  
 270 the number of recorded gargles/twargles per target population. “Gargle Hours”  
 271 and “Twargle Hours” refer to the amount of time, in hours, that gargle/twargle  
 272 data was collected by someone trained to collect data on the respective  
 273 vocalization. “Gargle Rate” and “Twargle Rate” refer to the target population’s  
 274 average gargle/twargle per hour rate.

275

	Gargle Count	Gargle Hours	Gargle Rate	Twargle Count	Twargle Hours	Twargle Rate
Infant (<8 mo.)	7080	901.16	7.86	409	359.73	1.14
Infant (<24 mo. including <8 mo.)	11763	3163.70	3.72	863	1107.08	0.78
Juvenile (≥24 & <60 mo.)	2076	3500.30	0.59	582	1809.72	0.32
Pregnant females	451	1115.99	0.40	507	1036.24	0.49
Females with young (<8 mo.) infants	1280	1010.75	1.27	746	946.44	0.79
Non-pregnant adult females (≥ 60 mo.) without young infants	867	2404.03	0.36	605	2234.96	0.27
All adult males (≥ 60 mo.)	253	5621.85	0.05	57	5239.29	0.01
All adults (≥ 60 mo.)	2851	10152.66	0.28	1915	9456.97	0.20
Overall population	16690	16816.65	0.99	3360	12373.78	0.27

276 Data were collected on 357 individuals; 221 individuals were  
 277 observed producing gargles or twargles, and 136 individuals were never  
 278 observed producing gargles or twargles (this does not indicate that 38%

35

36

279 of the population does not gargle or twargle). Researchers observed

280 16816.65 gargle-hours, 12373.78 twargle-hours, 16,690 gargle

281 instances, and 3,360 twargle instances

## 282 **(e) Data Cleaning**

283 Initial data cleaning and preparation were performed by querying  
284 the MySQL database housing the Lomas Barbudal Monkey Project's data.

285 Later stages of data cleaning, exploratory data analysis and inferential

286 statistics were conducted using Python 3.8.2 (<https://www.python.org/>)

287 and R 4.0.2 (<http://www.R-project.org/>).

288 Individuals born between 1994 and 2012 were genotyped using  
289 standard procedures employed by Dr. L. Vigilant's primate genetics lab  
290 at MPI-EVAN (see Godoy, Vigilant, & Perry, 2016; Muniz & Vigilant, 2008  
291 for details). 11 individuals, in addition to those who died before fecal  
292 sample collection, have not been genotyped. Infants (<24 months) with  
293 unknown fathers produced 1,222 out of 12,626 gargle and twargle  
294 instances. We did not include these instances in analyses for P.4, as  
295 paternity is relevant.

## 296 **(f) Statistical Analysis**

### 297 **(f.1.) Negative Binomial Models**

298 We addressed predictions (P1 - P4) using negative binomial (NB)  
299 generalized linear mixed models (GLMM) in the glmmTMB package in the  
300 R 4.0.2 statistical environment (<https://www.R-project.org/>). We used the

37

38

301 `fitdistrplus` and `logspline` packages, exploratory data analyses (Zuur,  
302 Ieno, & Elphick, 2010) and Akaike information criterion (AIC) to  
303 determine the best: a) distribution and b) models, of all considered  
304 models. NB distributions are reasonable for our data and account for  
305 gargle and twargle rate variance due to uneven focal sampling and  
306 proportionally more data collected during alpha male takeover events  
307 (Consul & Jain, 1973; Davis & Wu, 2009; Kaempfer, 1995; Lindén &  
308 Mäntyniemi, 2011; Lord, Guikema, & Geedipally, 2008). We fit NB  
309 models using two NB distributions (`nbinom1` and `nbinom2` in the  
310 `glmmTMB` package) and report results from `nbinom1` models, as these  
311 were better fitting models according to AIC. All models include caller  
312 identity as a random effect to account for idiosyncratic vocalization  
313 rates.

314 Gargle and twargle rates are dependent variables measured by  
315 vocalization count/individual/hour followed. Inconsistent follow time per  
316 monkey is accounted for by conditional probability and a log offset  
317 variable (Lindén & Mäntyniemi, 2011). Independent variables differ per  
318 prediction but always specify the target group (population subset) of  
319 interest. Reference groups are population subsets excluding target  
320 groups. Individuals are considered once or twice per prediction: a) in the  
321 target group, b) in the reference group or c) in the target group and the

39

40

322 reference group, if data was collected on that individual during different  
323 reproductive/life history stages.

324         We calculated incidence rate ratios (IRR) for each model. (The  
325 estimate is the log of the IRR.) IRRs represent the ratio of event  
326 outcomes over a given time period:  $IRR = 1$  indicates that the target and  
327 reference groups vocalize at the same rates,  $<1$  indicates that the  
328 reference group vocalizes at a lower rate, and  $>1$  indicates that the  
329 target group vocalizes at a higher rate. We present exponentiated  
330 confidence intervals (CI), as they relate to IRRs. If the CI includes 1, then  
331 there may not be a true difference between groups. P-values are two-  
332 tailed.

333         Table 1 outlines the target group and reference group for each  
334 analysis. We subcategorize infants into those aged 0-8 months and 8-23  
335 months. Infants were defined as  $<8$  months for all analyses except for  
336 P4a, as weaning occurs between 8-23 months of age (Carnegie, Fedigan,  
337 & Melin, 2011; Jack & Fedigan, 2008; Sargeant, Wikberg, Kawamura, &  
338 Fedigan, 2015). We used the lower-bound weaning age, because, if  
339 necessary, infants may have the ability to gain independence and avoid  
340 infanticide (Treves, 2000). Pregnant females are defined as females who  
341 will give birth in the subsequent 160 days. We define adults as  
342 individuals 60 months or older, because the youngest female to give  
343 birth in the population conceived at 60 months old (Perry, 2012).

41

42

**344 (f.2.) Binomial tests**

345 We use a binomial test to address this prediction, because we  
346 measure gargles and twargles as proportions, rather than  
347 rates/individual. Binomial tests compare two outcomes with the null  
348 assumption that each outcome will occur 50% of the time (Kaempf,  
349 1995). We compare gargles and twargles directed to the alpha male,  
350 versus to all other members of the group, from infants (<8 months),  
351 pregnant females, and female with infants (<8 months). We assume that  
352 there are at least two potential gargle or twargle recipients, which is  
353 true, as the smallest group at any given time was 7 individuals. An  
354 outcome over 50% indicates that monkeys gargle and twargle more to  
355 the alpha male than to all other group members combined.

**356 (f.3.) Data availability statement**

357 The data supporting the findings of this study are available in the  
358 supplementary materials of this article.

359

**360 Results****361 P.1: Infants will gargle and twargle at higher rates than the  
362 overall population.**

363 NB models suggest that infants gargle nearly eight times as much  
364 as the overall population:  $\sim 7.86$  gargle/hour compared to  $\sim .99$   
365 gargle/hour (NB:  $n=411$  observations,  $N=355$  groups,  $Z=23.91$ ,

43

44

366  $P < 0.0001$ ); see Table 2. Infants twargle 4.22 times as much as the  
367 overall population:  $\sim 1.14$  twargle/hour compared to  $\sim 0.27$  twargle/hour  
368 (NB:  $n = 400$  observations,  $N = 347$  groups,  $Z = 5.70$ ,  $P < 0.0001$ ; Figure 1;  
369 Tables 2 and 3).

370       Figure 1 (a plot of raw data) demonstrates the rate of change in  
371 gargle and twargle vocalizations with age. Monkeys gargle more than  
372 they twargle at almost all ages, exhibiting a general decline from four  
373 months of age onwards. The peak gargle and twargle rates occur during  
374 the 4<sup>th</sup> month of life, when infants are leaving their mothers' backs and  
375 begin exploring the world independently.

376

377

45

46

378 Table 3. Incident rate ratios (IRR), exponentiated confidence intervals (CI),  
 379 standard errors of the estimates (SE), random intercept variance (Var) and P-  
 380 values for the fixed effects for P.1: Infants will gargle and twargle at higher  
 381 rates than the overall population.  
 382

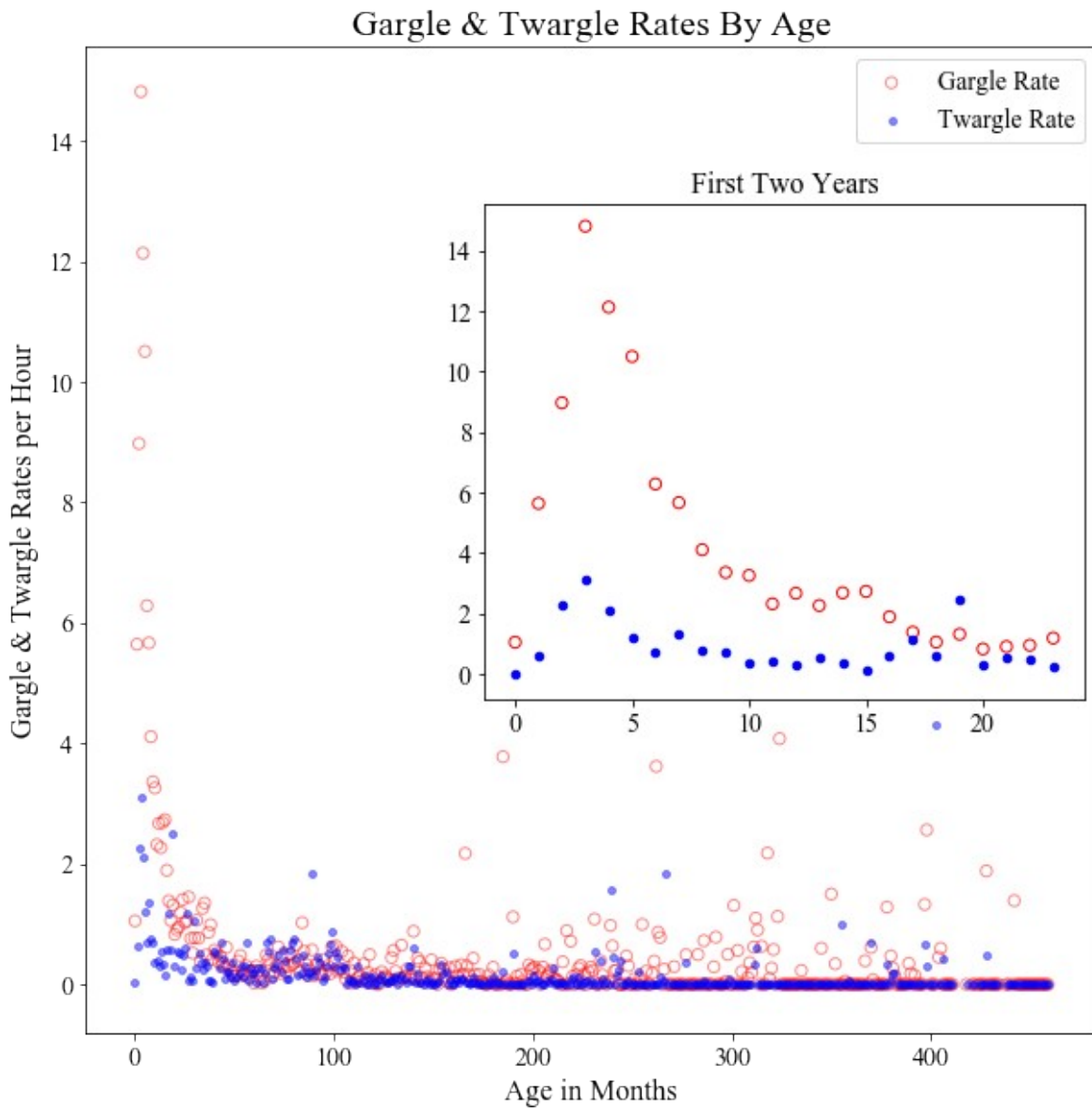
	IRR	CI	SE	Var	P-value	Prediction supported?
Gargle model of infants compared to the rest of the population	10.405	8.588 - 12.608	0.098	0.724	<0.0001	yes
Twargle Model of infants compared to the rest of the population	2.902	2.012 - 4.187	0.187	1.429	<0.0001	yes

383



47

48



384 Figure 1: The raw data scatterplot displays average gargle (open red circles)  
 385 and twargle (closed blue circles) rate data by age in months. The subplot  
 386 displays the average gargle and twargle rates for individuals 24 months of age  
 387 and younger.

388

389 **P.2: Pregnant females will gargle and twargle at higher rates**  
 390 **than the rest of the adult female population, especially adult**

49

50

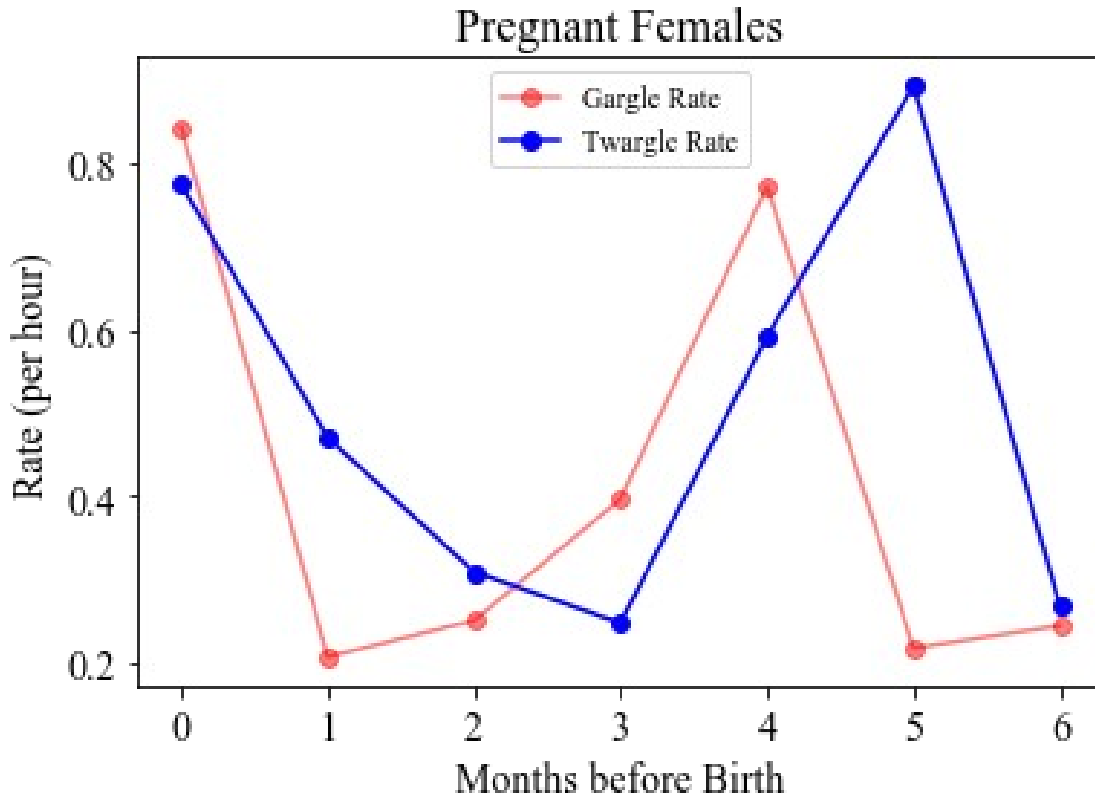
391 **females who are not pregnant and do not have an infant <8**  
 392 **months old.**

393 Females increase gargle and twargle rates during the birthing  
 394 month, as displayed by Figure 2 (a plot of raw data). Females show  
 395 marked increases in gargle and twargle rates around four and five  
 396 months before giving birth, respectively. We suspect the pattern is  
 397 related to pregnancy recognition and hormonal changes, but we do not  
 398 have data to test this theory.

399 Table 4. Incident rate ratios (IRR), exponentiated confidence intervals (CI),  
 400 standard errors of the estimates (SE), random intercept variance (Var) and P-  
 401 values for the fixed effects for P.2 models: Pregnant females will gargle and  
 402 twargle at higher rates than the rest of the adult female population, especially  
 403 adult females who are not pregnant and do not have an infant <8 months old.  
 404

	IRR	CI	SE	Var	P-value	Prediction supported?
Gargle model of pregnant females compared to non-pregnant adult females	0.717	0.535 - 0.961	0.150	0.312	0.026	no (wrong direction)
Twargle model of pregnant females compared to non-pregnant adult females	0.913	0.694 - 1.200	0.140	0.627	0.513	no
Gargle model of pregnant females compared to adult females who are not pregnant and do not have an infant <8 months old.	1.389	1.015 - 1.901	0.160	0.155	0.040	weak support
Twargle model of pregnant females compared to adult females who are not pregnant and do not have an infant <8 months old.	1.349	1.016 - 1.792	0.145	0.815	0.038	weak support

51  
52  
405



406 Figure 2: The raw data line graph displays the average gargle rates (red) and  
 407 twargle rates (blue) of pregnant females 0-6 months before giving birth. The  
 408 graph shows increases in gargle and twargle rates during the birthing month  
 409 and four and five months before giving birth, respectively.

410

411 P.2.a. Results suggest that pregnant females do not gargle and  
 412 twargle more than adult females who are not pregnant. (Gargle NB:  
 413 n=208 observations, N=112 groups, Z=-2.23, P=0.026; Twargle NB:  
 414 n=202 observations, N=111 groups, Z=-0.65, P=0.513).

415 P.2.b. Results suggest that pregnant females do not gargle and  
 416 twargle more than adult females who are not pregnant and do not have

53

54

417 an infant <8 months old (Gargle NB: n=206 observations, N=110  
418 groups, Z=2.05, P=0.040; Twargle NB: n=201 observations, N=110  
419 groups, Z=2.073, P=0.038; Table 4; Figure S1 - Figure S2).

420 **P.3: Females with infants (<8 months) will gargle and twargle at**  
421 **higher rates than the rest of the adult female population,**  
422 **especially adult females who are not pregnant and do not have**  
423 **an infant <8 months old.**

424 P.3.a. Results suggest that females with infants (<8 months)  
425 gargle and twargle more than the rest of the adult female population  
426 (Gargle NB: n=210 observations, N=112 groups, Z=9.09, P<0.0001;  
427 Twargle NB: n=206 observations, N=111 groups, Z=5.62, P<0.0001).

428 P.3.b. Results suggest that females with infants (<8 months)  
429 gargle and twargle more than adult females who are not pregnant and  
430 do not have an infant <8 months old (Gargle NB: n =210 observations,  
431 N=112 groups, Z=8.46, P<0.0001; Twargle NB: n=205 observations,  
432 N=110 groups, Z=6.21, P<0.0001; Table 5; Figures S3 - Figure S4).

433

55

56

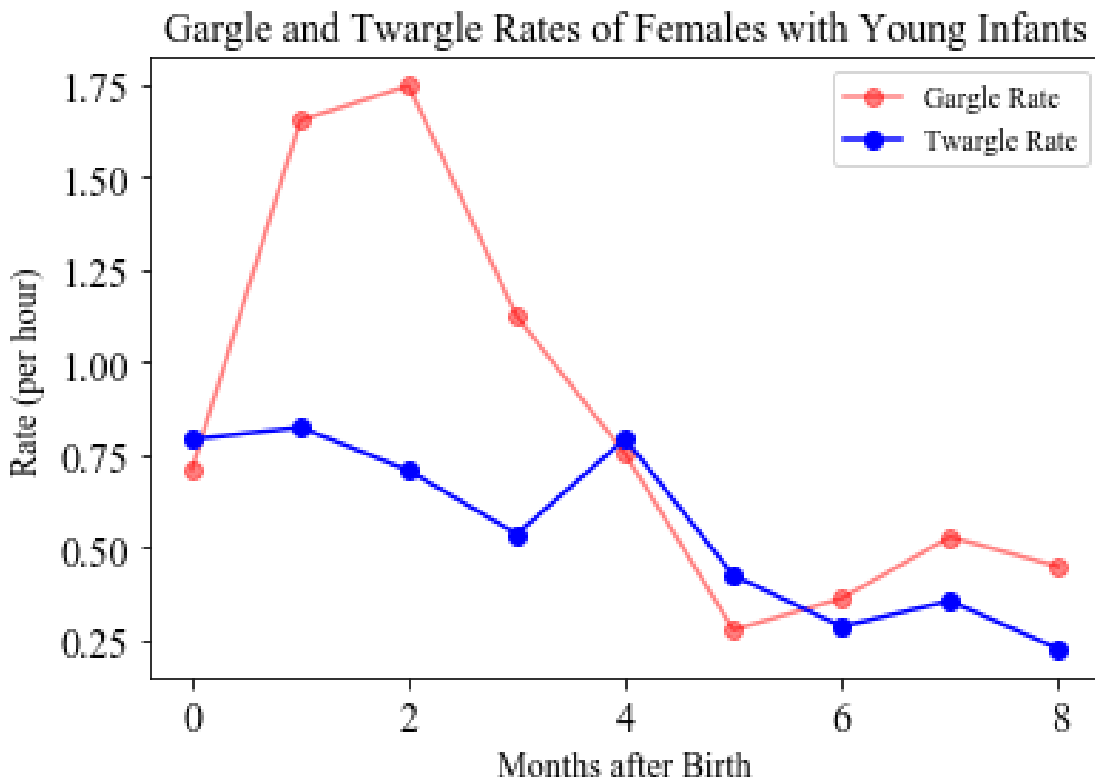
434 Table 5. Incident rate ratios (IRR), exponentiated confidence intervals (CI),  
 435 standard errors of the estimates (SE), random intercept variance (Var) and P-  
 436 values for the fixed effects for P.3 models: Females with infants (<8 months)  
 437 will gargle and twargle at higher rates than the rest of the adult female  
 438 population, especially adult females who are not pregnant and do not have an  
 439 infant <8 months old.

440

	IRR	CI	SE	Var	P-value	Prediction supported?
Gargle model of females with infants (<8 months) compared to the rest of adult females	3.086	2.421 - 3.935	0.124	0.237	<0.0001	yes
Twargle model of females with infants (<8 months) compared to adult females who are not pregnant and do not have an infant <8 months old.	2.087	1.615 - 2.698	0.131	0.582	<0.0001	yes
Gargle model of females with infants (<8 months) compared to adult females who are not pregnant and do not have an infant <8 months old.	3.416	2.570 - 4.541	0.145	0.201	<0.0001	yes
Twargle model of females with infants (<8 months) compared to adult females who are not pregnant and do not have an infant <8 months old.	2.341	1.790 - 3.061	0.137	0.645	<0.0001	yes

57  
 58  
 441  
 442  
 443  
 444  
 445

Females with infants (<8 months) gargle the most 1-2 months after giving birth and twargle the most 4 months after giving birth (Figure 3). They gargle more than they twargle during the first 3 months post-partum.



446 Figure 3: The raw data line graph displays the average gargle rates (red) and  
 447 twargle rates (blue) of females with young infants (<8 months). The graph  
 448 suggests that females with infants (<8 months) gargle at high rates 1-2  
 449 months after giving birth and at slightly higher rates 7-8 months after giving  
 450 birth. Females twargle at high rates 1-4 months after giving birth.  
 451 An alternative modeling approach to answering P2 and P3 is  
 452 presented in the SI (Table S5).

59

60

453 **P.4: Individuals will gargle and twargle more when the alpha**  
454 **male is not the offspring's (or fetus') father.**

455 P.4.a. Our results suggest that infants (<8 months) do not gargle  
456 or twargle more when the alpha male is not their father (Gargle NB:  
457 n=42 observations, N=40 groups, Z=0.173, P=0.863; Twargle NB: n=36  
458 observations, N=33 groups, Z=0.334, P=0.738). However, infants (8-23  
459 months) gargle, but may not twargle, more when the alpha male is not  
460 their father (Gargle NB: n=63 observations, N=48 groups, Z=3.436,  
461 P<0.001; Twargle NB: n=49 observations, N=45 groups, Z=1.880,  
462 P=0.060).

463 P.4.b. Our results suggest that pregnant females do not gargle or  
464 twargle more when the alpha male is not their father (Gargle NB: n=61  
465 observations, N=51 groups, Z=0.223, P=0.824; Twargle NB: n=55  
466 observations, N=46 groups, Z=-3.24, P=0.001).

467 P.4.c. Our results suggest that females with infants (<8 months)  
468 do not gargle or twargle more when the alpha male is not their father  
469 (Gargle NB: n=73 observations, N=58 groups, Z=0.433, P=0.665;  
470 Twargle NB: n=67 observations, N=53 groups, Z=-1.80, P=0.072; Table  
471 6).

472

473 Table 6. Incident rate ratios (IRR), exponentiated confidence intervals (CI),  
474 standard errors of the estimates (SE), random intercept variance (Var) and P-

61

62

475 values for the fixed effects for P.4 models: Individuals will gargle and twargle

476 more when the alpha male is not the offspring's (or fetus') father.

	IRR	CI	SE	Var	P-value	Prediction Supported?
Gargle model of infants (<8 months) whose father is not the alpha male compared to infants (<8 months) whose father is the alpha male	1.043	0.644 - 1.690	0.246	0.101	0.863	no
Twargle model of infants (<8 months) whose father is not the alpha male compared to infants (<8 months) whose father is the alpha male	1.211	0.394 - 3.723	0.573	0.339	0.738	no
Gargle model of infants (8 - 23 months) whose father is not the alpha male compared to infants (8 - 23 months) whose father is the alpha male	2.235	1.413 - 3.536	0.234	0.470	<0.001	yes
Twargle model of infants (8 - 23 months) whose father is not the alpha male compared to infants (8 - 23 months) whose father is the alpha male	1.914	0.973 - 3.766	0.345	0.598	0.060	no
Gargle model of pregnant females whose fetus' father is not the alpha male compared to pregnant females whose expected offspring's father is the alpha male	1.092	0.503 - 2.373	0.396	0.283	0.824	no
Twargle model of pregnant females whose fetus' father is not the alpha male compared to pregnant females whose expected offspring's father is the	0.350	0.185 - 0.660	0.324	1.486	0.001	no (wrong direction)



63

64

alpha male						
Gargle model of females with infants (<8 months) whose offspring's father is not the alpha male compared to females with infants (<8 months) whose offspring's father is the alpha male	1.1 58	0.597 - 2.244	0.338	0.374	0.66 5	no
Twargle model of females with infants (<8 months) whose offspring's father is not the alpha male compared to females with infants (<8 months) whose offspring's father is the alpha male	0.6 46	0.402 - 1.040	0.243	1.548	0.07 2	no

477

478

65

66

479 **P.5: Individuals will gargle and twargle more to the alpha male**  
480 **than to any other monkey in the group.**

481       Individuals gargled and twargled more to the alpha male than to  
482 all other individuals combined (binomial tests: both  $P < 0.0001$ ; see tables  
483 Table S2, S3 and S4 for further details) despite multiple subordinate  
484 males and only one alpha male occupying a group at any point in time.  
485 Throughout the entire study population, alpha males received 7628  
486 (57.9%) of the 13185 gargles produced, and 2243 (82.2%) of the 2729  
487 twargles produced. Infants (<8 months) twargled more to the alpha  
488 male, but they did not gargle more to the alpha male; 46% of gargles  
489 were directed at the alpha male (see SI Tables 2-4 for further  
490 information).

#### 491 **Discussion**

492       Our research explores our proposed, “Assessment of Infanticide  
493 Risk Hypothesis,” suggesting that white-faced capuchins use two  
494 elements of their vocal repertoire, the gargle and the twargle, to perform  
495 Zahavian bond tests. We predicted that individuals will gargle and  
496 twargle more frequently when facing infanticide risk, and we predicted  
497 that individuals will gargle and twargle more frequently to individuals  
498 posing infanticide risk, namely the alpha male. Our results provide some  
499 support for our hypothesis, but a few questions remain unclear.  
500 Individuals facing infanticide risk (to themselves or their offspring)

67

68

501 gargled and twargled at the highest rates out of all target groups in the  
502 study. However, pregnant females, whose fetuses were at risk for future  
503 infanticide, did not gargle and twargle at high rates. Furthermore,  
504 infants (8-23 months) were the only demographic group included in the  
505 study that gargled more to the alpha male when he was not their father.  
506 Infants (<8 months) were also the only demographic group included in  
507 the study that did not gargle more to the alpha male than to any other  
508 group member.

509        Infants (<8 months) gargle and twargle more than any other  
510 demographic group included in our study (Figure 1, Table 2). However,  
511 infants do not gargle or twargle more to the alpha male when he is not  
512 their father, and they do not gargle more to the alpha male than to any  
513 other group member. Our results suggest that assessing infanticide risk  
514 is not the sole motivation for infant gargling and twargling. One potential  
515 explanation is that infants may be strongly motivated to test bonds with  
516 multiple group members, because they may not have prior knowledge  
517 regarding allies and enemies. Infants (8-23 months) gargle more when  
518 the alpha male is not their father, and this could potentially represent  
519 the age at which individuals begin developing a sense of the social  
520 hierarchy. Also, this is the age in which infants begin spending less time  
521 around their mothers (Perry et al., 2012), and they may recognize the  
522 necessity of testing bonds with the alpha male when they receive less

69

70

523 protection from their mothers. They could also be motivated to test  
524 bonds to: 1) assess coalitionary support or 2) assess access to valuable  
525 resources.

526         Our study does not provide support for our prediction that  
527 pregnant females gargle and twargle at especially high rates (Table 4).  
528 We suspect that pregnant females, as compared to females with infants  
529 (<8 months), experience less selective pressure to test bonds, as their  
530 threat of infanticide is not so imminent as that of females with infants.

531         Adult females with infants (<8 months) gargle and twargle at high  
532 rates, especially when infants are young (<3 months old) (Figure 3).  
533 Mothers may vocalize at high rates during this time to convey gargle and  
534 twargle function and context to offspring, as researchers have found that  
535 mothers across many non-human primate species convey vocal context  
536 and function (Bergman et al., 2019; Elowson et al., 1998; Seyfarth &  
537 Cheney, 1986). Future analyses should investigate gargles and twargles  
538 in the context of mother-offspring dyads to address this possibility and  
539 the relationship between weaning and gargles and twargles.

540         Our results suggest that, with the exception of infants (8-23)  
541 months, individuals do not gargle nor twargle more when the alpha male  
542 is not their father (Table 6). We expected increased bond testing when  
543 the alpha male did not sire the offspring, as these are expected to be  
544 times when individuals are particularly at risk of infanticide and/or in

71

72

545 need of coalitionary support and resources (Perry, 2012). However, most  
546 of our studied demographic groups test bonds regardless of who holds  
547 the current alpha male position. We suggest several potential  
548 explanations: a) Individuals are motivated to test bonds, because they  
549 need coalitionary support and access to resources in many situations  
550 (e.g. during fights or inter-group encounters), b) Bond testing indicates  
551 long-term sentiments of a relationship that are likely to carry forward  
552 well into the future (Zahavi, 1977b; Zahavi & Zahavi, 1977), so  
553 individuals may preemptively test bonds in preparation of a risky  
554 situation, or c) Individuals may not recognize their father or trust that  
555 their father recognizes them as kin. However, according to the logic of  
556 (b), we would expect pregnant females to gargle and twargle at high  
557 rates if individuals preemptively test bonds, which we did not find, so  
558 explanation (b) seems unlikely.

559         The majority of gargles and the overwhelming majority of twargles  
560 were directed to the alpha male, across almost all demographic groups.  
561 This finding supports our hypothesis that individuals use gargles and  
562 twargles to test important bonds. The difference in gargle and twargle  
563 rates throughout the study seems to indicate that gargles and twargles  
564 are at least sometimes used for distinct purposes. Elements of the  
565 benign trill vocalization are incorporated into the twargle (Gros-Louis,  
566 2002; Gros-Louis et al., 2008), so it is possible that twargles are

73

74

567 produced with more benign intent than gargles. This may explain the  
568 difference in gargle and twargle rates of females with infants (<8  
569 months). Females gargle, rather than twargle, at high rates 0-3 months  
570 post-partum (Figure 3). Perhaps this is an especially crucial time for  
571 females to bond test (Perry et al., 2012), because infants are  
572 nutritionally dependent and at highest risk for infanticide, which may  
573 influence vocalization selection.

574       Alternative hypotheses could explain gargle and twargle functions.  
575 We considered alternative hypotheses previously (Perry 1998; Perry &  
576 Manson, 2008) and they were not supported because of the context in  
577 which gargles and twargles are produced. If gargles were used to  
578 formally acknowledge another's superior rank, we would expect them to  
579 be frequent and unidirectional in most dyads of disparate rank and least  
580 frequent in adjacently ranked individuals during times when relative  
581 rank was being disputed. If the gargles were used to test bonds with  
582 adjacently ranked individuals, they should be more frequently  
583 exchanged (in both directions) within these dyads (Preuschoft, 1999;  
584 Smith et al., 2011). Neither of these situations is true, as gargles are  
585 primarily used by infants and their mothers towards adult (and primarily  
586 alpha) males. If individuals use (gargle) vocalizations as appeasement,  
587 to mitigate conflict and tension, they would be expected to occur before,  
588 during or shortly after conflicts if (Dias, Luna, & Espinosa, 2008;

75

76

589 Hohmann & Fruth, 2000; Smith et al., 2011); however, gargles and  
590 twargles are produced primarily during peaceful interactions. If  
591 gargles/twargles are indicative of respect by the vocalizer towards the  
592 recipient, we would expect the recipient to respond neutrally or  
593 affiliatively; however, a negative response is often produced (Gros-Louis  
594 et al., 2008; Perry & Manson, 2008). And if gargles and twargles are a  
595 sign of allegiance, then alpha males should be concerned when they  
596 hear gargles being directed towards other group members, yet they pay  
597 no attention when these situations arise (Perry & Manson, 2008).

598         Therefore, “The Assessment of Infanticide Risk” remains our  
599 leading hypothesis explaining gargles and twargles.

#### 600 **(a) Limitations**

601         Although this study produces results consistent with the  
602 Assessment of Infanticide Risk hypothesis by comparing gargle and  
603 twargle rates across demographic categories, it would be desirable to  
604 also conduct a fine-grained temporal analysis of how relationships  
605 change within each dyad, looking at the causes and consequences of  
606 producing gargles and twargles. Are these vocalizations produced when  
607 the relationship is under particular strain? Is the vocalizer’s subsequent  
608 behavior contingent on the affective responses of the recipient to the  
609 gargle or twargle? The current data set is not well suited to this goal,  
610 because of the sampling density within each dyad. However, these

77

78

611 would be exciting analyses to conduct with a more limited sample of  
612 individuals that have dense focal sampling within time periods  
613 characterized by changes in relationship quality.

#### 614 **(b) Conclusion**

615 Our results largely support the Assessment of Infanticide Risk  
616 Hypothesis. Infants and females with infants (<8 months), i.e.  
617 individuals/their offspring at greatest infanticide risk, gargle and twargle  
618 at higher rates than the overall population and population subsets.  
619 Pregnant females did not gargle and twargle at exceptionally high rates,  
620 perhaps because their potential infanticide risk was too far in their  
621 future. Most demographic groups gargle and twargle more to the alpha  
622 male than to any other and all other individuals in the group. However,  
623 with the exception of infants 8-23 months of age, individuals do not  
624 gargle or twargle more when the alpha male is not their father. Overall,  
625 we found that individuals are motivated to test bonds during infanticidal  
626 risk periods, but our results suggest that that individuals may be  
627 motivated to test bonds for alternative reasons as well.

#### 628 **Funding**

629 The fieldwork component of this project was funded by the  
630 following grants to S. Perry: National Science Foundation (1638428,  
631 0613226, 848360, 1232371), the National Geographic Society (7968-06,  
632 8671-09, 20113909, 9795-15), the Templeton World Charity Foundation,



79

80

633 Inc. (grant 0208), and 4 grants from the L.S.B. Leakey Foundation.

634 Additional financial support was provided by the Max Planck Institute for

635 Evolutionary Anthropology, UCLA and the Wild Capuchin Foundation.

### 636 **Acknowledgments**

637 The following people assisted S. Perry and Alexa Duchesneau in

638 substantive behavioral data collection, data processing, and/or field sites

639 logistics for this project: C. Angyal, A. Autor, B. Barrett, L. Beaudrot, M.

640 Bergstrom, R. Berl, A. Bjorkman, L. Blankenship, T. Borcuch, J. Broesch,

641 D. Bush, J. Butler, F. Campos, C. Carlson, S. Caro, A. Cobden, M. Corrales,

642 J. Damm, C. Dillis, N. Donati, C. de Rango, A. Davis, G. Dower, R. Dower,

643 K. Feilen, J. Fenton, K. Fisher, A. Fuentes J., M. Fuentes, T. Fuentes A., C.

644 M. Gault, H. Gilkenson, I. Godoy, I. Gottlieb, J. Gričiute, L.M. Guevara R.,

645 L. Hack, R. Hammond, S. Herbert, C. Hirsch, M. Hoffman, A. Hofner, C.

646 Holman, J. Hubbard, S. Hyde, M. Jackson, O. Jacobson, E. Johnson, L.

647 Johnson, K. Kajokaite, M. Kay, E. Kennedy, D. Kerhoas-Essens, S. Kessler,

648 W. Krimmel, W. Lammers, S. Lee, S. Leinwand, S. Lopez, T. Lord, J.

649 Mackenzie, S. MacCarter, M. Mayer, F. McKibben, A. Mensing, W. Meno,

650 M. Milstein, C. Mitchell, Y. Namba, D. Negru, A. Neyer, C. O'Connell, J.C.

651 Ordoñez J., N. Parker, B. Pav, R. Popa, K. Potter, K. Ratliff, K. Reinhardt,

652 N. Roberts Buceta, E. Rothwell, H. Ruffler, S. Sanford, C. M. Saul, I.

653 Schamberg, N. Schleissman, C. Schmitt, S. Schulze, A. Scott, E.

654 Seabright, J. Shih, S. Sita, K. Stewart, K. van Atta, L. van Zuidam, J.

81

82

655 Vandermeer, J. Verge, V. Vonau, A. Walker-Bolton, E. Wikberg, E.  
656 Williams, E. Wolf, and D. Wood. We are particularly grateful to W.  
657 Lammers and H. Gilkenson, for long-term management of the field site  
658 from 2001-2013. We extend extreme gratitude to D. Cohen for data  
659 management assistance, and helpful discussion at all stages. Both D.  
660 Cohen and A. Lin provided helpful statistical advice.

661       Permission to conduct the research was provided by the Costa  
662 Rican Park Service (SINAC and ACAT), C. Jiménez Freer (of Brin d'Amor),  
663 El Pelón de la Bajura, and the residents of San Ramon de Bagaces. All  
664 opinions, findings and conclusions or recommendations expressed in this  
665 material are those of the authors and do not necessarily reflect the  
666 views of the National Science Foundation or other funding agencies.

## 667 **References**

- 668 Bergman, T. J., Beehner, J. C., Painter, M. C., & Gustison, M. L. (2019).  
669       The speech-like properties of nonhuman primate vocalizations.  
670       *Animal Behaviour*, *151*, 229–237.  
671       <https://doi.org/10.1016/j.anbehav.2019.02.015>
- 672 Carnegie, S. D., Fedigan, L. M., & Melin, A. D. (2011). Reproductive  
673       Seasonality in Female Capuchins (*Cebus capucinus*) in Santa Rosa  
674       (Area de Conservación Guanacaste), Costa Rica. *International Journal*  
675       *of Primatology*, *32*(5), 1076–1090. [https://doi.org/10.1007/s10764-](https://doi.org/10.1007/s10764-011-9523-x)  
676       011-9523-x

83

84

- 677 Cheney, D. L., & Seyfarth, R. M. (2018). Flexible usage and social  
678 function in primate vocalizations. *Proceedings of the National*  
679 *Academy of Sciences of the United States of America*, 115(9), 1974–  
680 1979. <https://doi.org/10.1073/pnas.1717572115>
- 681 Consul, P. C., & Jain, G. C. (1973). A generalization of the poisson  
682 distribution. *Technometrics*, 15(4), 791–799.  
683 <https://doi.org/10.1080/00401706.1973.10489112>
- 684 Davis, R. A., & Wu, R. (2009). A negative binomial model for time series  
685 of counts. *Biometrika*, 96(3), 735–749.  
686 <https://doi.org/10.1093/biomet/asp029>
- 687 Dias, P. A. D., Luna, E. R., & Espinosa, D. C. (2008). The functions of the  
688 “greeting ceremony” among male mantled howlers (*Alouatta*  
689 *palliata*) on Agaltepec Island, Mexico. *American Journal of*  
690 *Primatology*, 70(7), 621–628. <https://doi.org/10.1002/ajp.20535>
- 691 Elowson, M. A., Snowdon, C. T., & Lazaro-Perea, C. (1998). Infant ‘  
692 Babbling’ in a Nonhuman Primate: Complex Vocal Sequences with  
693 Repeated Call Types. *Behavior*, 135(5), 643–664.
- 694 Fedigan, L. M. (2003). Impact of male takeovers on infant deaths, births  
695 and conceptions in *Cebus capucinus* at Santa Rosa, Costa Rica.  
696 *International Journal of Primatology*, 24(4), 723–741.  
697 <https://doi.org/10.1023/A:1024620620454>
- 698 Frankie, G. W., Vinson, S. B., Newstrom, L. E., Barthell, J. F., Rican, C., &

85

86

699 Forest, D. (1988). Nest Site and Habitat Preferences of Centris Bees  
700 in the Costa Rican Dry Forest. *Biotropica*, 20(4), 301–310.

701 Godoy, I., Vigilant, L., & Perry, S. E. (2016). Cues to kinship and close  
702 relatedness during infancy in white-faced capuchin monkeys, *Cebus*  
703 *capucinus*. *Animal Behaviour*, 116, 139–151. [https://doi.org/10.1016/](https://doi.org/10.1016/j.anbehav.2016.03.031)  
704 [j.anbehav.2016.03.031](https://doi.org/10.1016/j.anbehav.2016.03.031)

705 Gros-Louis, J. J. (2002). Contexts and behavioral correlates of trill  
706 vocalizations in wild white-faced capuchin monkeys (*Cebus*  
707 *capucinus*). *American Journal of Primatology*, 57(4), 189–202. [https://](https://doi.org/10.1002/ajp.10042)  
708 [doi.org/10.1002/ajp.10042](https://doi.org/10.1002/ajp.10042)

709 Gros-Louis, J. J. (2006). Acoustic analysis and contextual description of  
710 food-associated calls in white-faced capuchin monkeys (*Cebus*  
711 *capucinus*). *International Journal of Primatology*, 27(1), 273–294.  
712 <https://doi.org/10.1007/s10764-005-9012-1>

713 Gros-Louis, J. J., Perry, S. E., Fichtel, C., Wikberg, E., Gilkenson, H.,  
714 Wofsy, S., & Fuentes, A. (2008). Vocal repertoire of *Cebus capucinus*:  
715 Acoustic structure, context, and usage. *International Journal of*  
716 *Primatology*, 29(3), 641–670. [https://doi.org/10.1007/s10764-008-](https://doi.org/10.1007/s10764-008-9263-8)  
717 [9263-8](https://doi.org/10.1007/s10764-008-9263-8)

718 Hohmann, G., & Fruth, B. (2000). Use and function of genital contacts  
719 among female bonobos. *Animal Behaviour*, 60(1), 107–120.  
720 <https://doi.org/10.1006/anbe.2000.1451>

87

88

- 721 Hrdy, S. B. (1979). Infanticide among animals: A review, classification,  
722 and examination of the implications for the reproductive strategies  
723 of females. *Ethology and Sociobiology*, 1, 13-40.
- 724 Jack, K. M., & Fedigan, L. M. (2004). Male dispersal patterns in white-  
725 faced capuchins, *Cebus capucinus* Part 2: Patterns and causes of  
726 secondary dispersal. *Animal Behaviour*, 67(4), 771-782.  
727 <https://doi.org/10.1016/j.anbehav.2003.06.015>
- 728 Jack, K. M., & Fedigan, L. M. (2008). Female Dispersal in a Female-  
729 Philopatric Species , *Cebus capucinus*. *Behaviour*, 146(4), 471-497.
- 730 Kaempf, U. (1995). The Binomial Test: A Simple Tool to Identify Process  
731 Problems. *IEEE Transactions on Semiconductor Manufacturing*, 8(2),  
732 160-166. <https://doi.org/10.1109/66.382280>
- 733 Lindén, A., & Mäntyniemi, S. (2011). Using the negative binomial  
734 distribution to model overdispersion in ecological count data.  
735 *Ecology*, 92(7), 1414-1421. <https://doi.org/10.1890/10-1831.1>
- 736 Locke, J. L. (2001). Rank and relationships in the evolution of spoken  
737 language. *Journal of the Royal Anthropological Institute*, 7(1), 37-50.  
738 <https://doi.org/10.1111/1467-9655.00049>
- 739 Lord, D., Guikema, S. D., & Geedipally, S. R. (2008). Application of the  
740 Conway-Maxwell-Poisson generalized linear model for analyzing  
741 motor vehicle crashes. *Accident Analysis and Prevention*, 40(3),  
742 1123-1134. <https://doi.org/10.1016/j.aap.2007.12.003>

89

90

- 743 Manson, J. (1999). Infant handling in wild *Cebus capucinus*: Testing  
744 bonds between females? *Animal Behaviour*, 57(4), 911–921.  
745 <https://doi.org/10.1006/anbe.1998.1052>
- 746 Maynard Smith, J. (1982). Do animals convey information about their  
747 intentions? *Journal of Theoretical Biology*, 97(1), 1–5. [https://doi.org/10.1016/0022-5193\(82\)90271-5](https://doi.org/10.1016/0022-5193(82)90271-5)
- 749 Muniz, L., & Vigilant, L. (2008). Isolation and characterization of  
750 microsatellite markers in the white-faced capuchin monkey (*Cebus*  
751 *capucinus*) and cross-species amplification in other New World  
752 monkeys. *Molecular Ecology Resources*, 8(2), 402–405.  
753 <https://doi.org/10.1111/j.1471-8286.2007.01971.x>
- 754 Palombit, R. A. (1999). Infanticide and the evolution of pair bonds in  
755 nonhuman primates. *Evolutionary Anthropology*, 7(4), 117–129.  
756 [https://doi.org/10.1002/\(SICI\)1520-6505\(1999\)7:4<117::AID-](https://doi.org/10.1002/(SICI)1520-6505(1999)7:4<117::AID-EVAN2>3.0.CO;2-O)  
757 [EVAN2>3.0.CO;2-O](https://doi.org/10.1002/(SICI)1520-6505(1999)7:4<117::AID-EVAN2>3.0.CO;2-O)
- 758 Perry, S. E. (1997). Male-female social relationships in wild white-faced  
759 capuchins (*Cebus capucinus*). *Behaviour*, 134(7–8), 477–510. <https://doi.org/10.1163/156853997X00494>
- 761 Perry, S. E. (1998). A case report of a male rank reversal in a group of  
762 wild white-faced capuchins (*Cebus capucinus*). *Primates*, 39(1), 51–  
763 70. <https://doi.org/10.1007/BF02557743>
- 764 Perry, S. E. (2011). Social traditions and social learning in capuchin

91

92

765 monkeys (*Cebus*). *Philosophical Transactions of the Royal Society B:*

766 *Biological Sciences*, 366(1567), 988–996.

767 <https://doi.org/10.1098/rstb.2010.0317>

768 Perry, S. E. (2012). The Behavior of Wild White-Faced Capuchins.

769 Demography, Life History, Social Relationships, and Communication.

770 In *Advances in the Study of Behavior* (Vol. 44).

771 <https://doi.org/10.1016/B978-0-12-394288-3.00004-6>

772 Perry, S. E., Baker, M., Fedigan, L. M., Gros-Louis, J. J., Jack, K. M.,

773 MacKinnon, K. C., ... Rose, L. (2003). Social conventions in wild

774 white-faced capuchin monkeys: Evidence for traditions in a

775 neotropical primate. *Current Anthropology*, 44(2), 241–268.

776 <https://doi.org/10.1086/345825>

777 Perry, S. E., Godoy, I., & Lammers, W. (2012). The Lomas Barbudal

778 Monkey Project: Two Decades of Research on *Cebus capucinus*. In P.

779 M. Kappeler & D. Watts (Eds.), *Long-Term Field Studies of Primates*

780 (pp. 141–164). <https://doi.org/10.1017/CBO9781107415324.004>

781 Perry, S. E., Godoy, I., Lammers, W., & Lin, A. (2017). Impact of

782 personality traits and early life experience on timing of emigration

783 and rise to alpha male status for wild male white-faced capuchin

784 monkeys (*Cebus capucinus*) at Lomas Barbudal Biological Reserve,

785 Costa Rica. *Behaviour*, 154(2), 195–226.

786 <https://doi.org/10.1163/1568539X-00003418>

93

94

- 787 Perry, S. E., & Manson, J. (2008). Manipulative Monkeys: The Capuchins  
788 of Lomas Barbudal. In *Block Caving – A Viable Alternative?* (Vol. 21).  
789 Cambridge, MA: Harvard University Press.
- 790 Perry, S. E., & Smolla, M. (2020). Capuchin monkey rituals: an  
791 interdisciplinary study of form and function. *BioRxiv*, 375(1805),  
792 2020.02.21.958223. <https://doi.org/10.1101/2020.02.21.958223>
- 793 Poole, J. H. (1989). Announcing intent: the aggressive state of musth in  
794 African elephants. *Animal Behaviour*, 37(PART 1), 153-155.  
795 [https://doi.org/10.1016/0003-3472\(89\)90015-8](https://doi.org/10.1016/0003-3472(89)90015-8)
- 796 Preuschoft, S. (1999). Are Primates Behaviorists.pdf. *Journal of*  
797 *Comparative Psychology*, Vol. 113, pp. 91-95.
- 798 Sargeant, E. J., Wikberg, E. C., Kawamura, S., & Fedigan, L. M. (2015).  
799 Allonursing in white-faced capuchins (*Cebus capucinus*) provides  
800 evidence for cooperative care of infants. *Behaviour*, 152(12-13),  
801 1841-1869. <https://doi.org/10.1163/1568539X-00003308>
- 802 Schamberg, I., Cheney, D. L., Clay, Z., Hohmann, G., & Seyfarth, R. M.  
803 (2017). Bonobos use call combinations to facilitate inter-party travel  
804 recruitment. *Behavioral Ecology and Sociobiology*, 71(4).  
805 <https://doi.org/10.1007/s00265-017-2301-9>
- 806 Seyfarth, R. M., & Cheney, D. L. (1986). Vocal development in vervet  
807 monkeys. *Animal Behaviour*, 34(6), 1640-1658.  
808 [https://doi.org/10.1016/S0003-3472\(86\)80252-4](https://doi.org/10.1016/S0003-3472(86)80252-4)



95

96

809 Silk, J. B., Kaldor, E., &amp; Boyd, R. (2000). Cheap talk when interests

810 conflict. *Animal Behaviour*, 59(2), 423–432.811 <https://doi.org/10.1006/anbe.1999.1312>

812 Silk, J. B., Seyfarth, R. M., &amp; Cheney, D. L. (2016). Strategic use of

813 affiliative vocalizations by wild female baboons. *PLoS ONE*, 11(10),814 1–10. <https://doi.org/10.1371/journal.pone.0163978>

815 Smith, J. E., Powning, K. S., Dawes, S. E., Estrada, J. R., Hopper, A. L.,

816 Piotrowski, S. L., &amp; Holekamp, K. E. (2011). Greetings promote

817 cooperation and reinforce social bonds among spotted hyaenas.

818 *Animal Behaviour*, 81(2), 401–415.819 <https://doi.org/10.1016/j.anbehav.2010.11.007>

820 Smuts, B., &amp; Watanabe, J. M. (1990). Social relationships and ritualized

821 greetings in adult male baboons (*Papio cynocephalus anubis*).822 *International Journal of Primatology*, 11(2), 147–172.823 <https://doi.org/10.1007/BF02192786>824 Taylor, D. J. (2014). *Evolution of the Social Contract submitted by*825 *University of Bath*.826 Tomasello, M. (1990). Chimpanzee Culture?\*. *National Student Speech*827 *Language Hearing Association Journal*, 18, 73–75.

828 Treves, A. (2000). Prevention of infanticide: the perspective of infant

829 primates. In C. P. van Schaik & C. H. Janson (Eds.), *Infanticide by*830 *Males and its Implications* (pp. 223–228). Cambridge University

97

98

831 Press.

832 Van Schaik, C. P., &amp; Dunbar, I. M. (1990). The Evolution of Monogamy in

833 Large Primates : A New Hypothesis and Some Crucial Tests.

834 *Behaviour*, 115(1), 30-62.

835 Zahavi, A. (1977a). Reliability in communication systems and the

836 evolution of altruism. In P. C. (Ed.), *Evolutionary Ecology* (pp. 253-

837 254). Palgrave, London: Stonehouse B.

838 Zahavi, A. (1977b). The Testing of a Bond. *Animal Behaviour*, 25(1), 246-

839 247.

840 Zahavi, A., & Zahavi, A. (1977). *The Handicap Principle: A Missing Piece*841 *of Darwin's Puzzle*. Oxford University Press.

842 Zuur, A. F., Ieno, E. N., &amp; Elphick, C. S. (2010). A protocol for data

843 exploration to avoid common statistical problems. *Methods in*844 *Ecology and Evolution*, 1(1), 3-14. <https://doi.org/10.1111/j.2041->

845 210x.2009.00001.x

846

99

100

847 Title: Are Demographic Correlates of White-faced Capuchin Monkeys (*Cebus*  
 848 *capucinus*) “Gargle and Twargle” Vocalization Rates Consistent with the  
 849 Infanticide Risk Assessment Hypothesis?

850

851

Supporting Information

852

853 **Characteristics of the Data Set**

854 Table S1 Distribution of behavioral sampling effort across demographic categories

Target Population	Estimated Percentage of Total Population	Percentage of Gargle Hours	Percentage of Twargle Hours
Infants (< 24 months)	21%	19%	9%
Juveniles (24 - 59 months)	21%	21%	15%
Adult Males (>= 60 months)	26%	33%	42%
Adult Females (>= 60 months)	32%	27%	34%

855 To estimate % of total population we do the following: Each monkey that has  
 856 been identified has been assigned an estimated birth date, which is used to  
 857 compute its (estimated) age on every date when it is seen. We count a  
 858 monkey as having been seen during a month of its life (e.g. at the age of 21  
 859 months) if that was its estimated age (discarding fractional months) any of  
 860 the times it was seen. The fraction of monkeys estimated to be some age (in  
 861 months) is the number of monkeys seen at that age divided by the sum of  
 862 the numbers seen for all ages. Clearly a monkey who was seen at age 9  
 863 months and then again at 11 months was alive at 10 months, but is not  
 864 counted.

865 The latter two columns are computed on the basis of the time spent  
 866 collecting focal follows of monkeys observed by researchers who could  
 867 reliably identify the specified vocalization.

868

869 **Additional Results**

870 **Distribution of gargles and twargles across age-sex categories for**  
 871 **vocalizers and targets of vocalizations:**

872 Tables S2 and S3 show the distribution of the sample according to the age-sex class  
 873 of vocalizers and their recipients. Note that these are raw frequencies, not rates of  
 874 behavior, i.e. the amount of focal follow time is not included here, so the  
 875 percentages in columns 2-5 are for the entire pooled sample, not for the percentage  
 876 directed specifically by that age-sex class to another age-sex. Focal animals are

101

102

877 always the vocalizers in our analyses. Therefore, in the final column, we present the  
 878 percentage of all vocalizations for this age-sex class that are directed towards adult  
 879 males, defined as being >5 years of age. We also define adult females as those  
 880 being >5 years of age. Except for adult males, all other age-sex classes direct their  
 881 gargles and twargles overwhelmingly towards adult males.

882

883 Table S2: Number of gargles in the sample, according to age-sex class of vocalizer  
 884 and age-sex class of recipient, for all cases in which age-sex classes of both are  
 885 known, followed by the corresponding % of all vocalizations in the entire sample in  
 886 parentheses. The final column is the % of vocalizations by that particular age-sex  
 887 class that are directed towards adult males.

888

Age-sex class of vocalizer	Age-sex class of recipient				% of gargles directed towards males (>5 years of age) for the specified age-sex class of vocalizer
	Females (<5 years of age)	Females (>5 years of age)	Males (< 5 years of age)	Males (>5 years of age)	
Females (≤5 years of age)	35 (0.24%)	50 (0.34%)	16 (0.11%)	6636 (45.73%)	98.50%
Females (>5 years of age)	7 (0.05%)	2 (0.01%)	5 (0.03%)	2297 (15.83%)	99.39%
Males (≤5 years of age)	19 (0.13%)	334 (2.30%)	55 (0.38%)	4966 (34.22%)	92.41%
Males (>5 years of age)	23 (0.16%)	23 (0.16%)	33 (0.23%)	33 (0.23%)	29.46%
Total	84 (0.58%)	409 (2.82%)	109 (0.75%)	13932 (96.00%)	

889

890

103  
104  
891  
892  
893  
894  
895  
896

Table S3: Number of twargles in the sample, according to age-sex class of vocalizer and age-sex class of recipient, for all cases in which age-sex classes of both are known, followed by the corresponding % of all vocalizations in the entire sample in parentheses. The final column is the % of vocalizations by that particular age-sex class that are directed towards adult males.

Age-sex class of vocalizer	Age-sex class of recipient				% of twargles directed towards males (>5 years of age) for the specified age-sex class of vocalizer
	Females (< 5 years of age)	Females (>5 years of age)	Males (< 5 years of age)	Males (>5 years of age)	
Females (< 5 years of age)	17 (0.61%)	10 (0.36%)	5 (0.18%)	1003 (36.20%)	96.91%
Females (>5 years of age)	17 (0.61%)	10 (0.36%)	34 (1.23%)	1375 (49.62%)	95.75%
Males (<5 years of age)	0 (0.00%)	5 (0.18%)	7 (0.25%)	245 (8.84%)	95.33%
Males (>5 years of age)	7 (0.25%)	9 (0.32%)	12 (0.43%)	15 (0.54%)	34.88%
Total	41 (1.48%)	34 (1.23%)	58 (2.09%)	2638 (95.20%)	

897  
898  
899  
900  
901  
902  
903  
904  
905  
906  
907  
908  
909  
910  
911  
912  
913  
914

**P.1: Infants will gargle and twargle at higher rates than the overall population.**

Fifty-six monkeys were observed both as infants and non-infants, permitting a more longitudinal approach than was used in the main text. These 56 individuals gargled at 8.23/hour when <8 months old and 0.96/hour when >8 months old. Forty-five gargled at higher rates during infancy as compared to adulthood. The 11 exceptions are mostly accounted for by small observation times as infants. One individual was observed for 9, one for 4.5 hrs, one for 2.2 hrs and the rest for < 2 hours. Only 2 of these 11 exceptions were observed gargling as infants. Twenty-seven monkeys were observed for > 10 hours as infants (and also for > 10 hours as non-infants), and all of these individuals gargled at higher average rates as infants than as non-infants.

Twargles are less commonly observed than gargles, so more observation time is required to obtain a similar sample size to the gargle data. Fifty-three individuals were observed (by observers competent to identify twargles) both as infants and non-infants, and only 8 were observed for at least ten hours as infants (and >100

105

106

915 hours as non-infants.) One individual twargled at a higher average rate as a non-  
 916 infant than as an infant, and the other 7 displayed higher rates as infants. In total,  
 917 only 20 of the 53 displayed higher average twargle rates as infants, but none of the  
 918 others (N=33) were observed twargling as infants (i.e. an average rate of zero).  
 919 Twelve individuals also displayed zero rates as non-infants. Eleven individuals were  
 920 observed for over 100 hours as non-infants, all of whom were observed twargling at  
 921 least once.

922 **P.5: Individuals will gargle and twargle more to the alpha male than to any**  
 923 **other monkey in the group.**

924 Table S4: Percentage of vocalizations directed towards alpha males as opposed to  
 925 all other group members combined.

926

Vocalization type	Percentage of Vocalizations directed to alpha male	Number of vocalizations directed to the alpha male	Number of vocalizations directed to all other individuals	P-value
Gargles	57.9%	7628	5557	<0.0001
Twargles	82.2%	2243	486	<0.0001

927

928 Table S5: Counts and percentages of gargles and twargles directed by infants to (a)  
 929 the alpha male, or (b) all other individuals in the group.

930

Behavior	To Alpha	Not to Alpha	Percentage to Alpha
Infant (<8 months) gargles	2611	3061	46%
Infant (<8 months) twargles	259	105	71%
Infant (8-23 months) gargles	1611	1697	49%
Infant (8-23 months) twargles	275	116	70%

931

932 Table S6: Counts and percentages of gargles and twargles directed by adults (>60  
 933 months old) to (a) the alpha male, or (b) all other individuals in the group.

934

Behavior	To	Not to	Percentage to Alpha
----------	----	--------	---------------------

107

108

	Alpha	Alpha	
Adult (>60 months) gargles	2006	348	85%
Adult (>60 months) twargles	1283	192	87%

935

936

109

110

**937 Interactions between Group Size and Gargle and Twargle Rates**

938 In response to a request from reviewers, we explore how group size and age of  
939 vocalizer might affect the proportion of gargles and twargles directed towards the  
940 alpha male, using Generalized Linear Mixed Model, using the glmmTMB function in  
941 R (<http://www.R-project.org/>) (results below in S7). Age and group size are fixed  
942 effects, and monkey id is a random effect:

943

944  $\text{alpha} \sim \text{age5} + \text{size} + \text{offset}(\log(\text{total})) + (1 \mid \text{id})$

945

946 “Alpha” is the fraction of gargles or twargles directed towards the alpha male,  
947 “age5” is the age category (as described below), “size” is the group size, “total” is  
948 the number of gargles or twargles produced, and “id” is the identity of an individual  
949 monkey. In the model, individuals are grouped into 6 age categories: each of the  
950 first 5 years of life is treated as a separate category (age 0 to 4) and the 6<sup>th</sup>  
951 category lumps all individuals age 5 years or more (i.e. adults). Group size ranges  
952 from 5 to 41 individuals. We ran models using the nbinom1 and nbinom2  
953 distributions, but we only report the nbinom2 results, as models using this  
954 distribution performed better according to AIC (Table S7). See the code and output  
955 file for further details.

956

957 The models show how age and group size are related to the fraction of gargles and  
958 twargles directed towards the alpha male. For every year of age, up to age 5, there  
959 is an increase by a factor of 1.094 in the fraction of gargles, and an increase by a  
960 factor of 1.032 in the fraction of twargles, directed to the alpha male. Also, for each  
961 individual added to the group, the fraction of gargles directed to the alpha male  
962 changes by a factor of 0.985 (i.e. decreases), and the fraction of twargles directed  
963 to the alpha male changes by a factor of 0.994 (also decreasing).

964

965 Summaries of the raw data showing how the proportion of gargles directed to the  
966 alpha male vary according to age and group size are presented after the model  
967 results, in Tables S8 & S9.

968



111

112

969 Table S7: Results from the GLMMs predicting the impact of (a) age of vocalizer and  
 970 (b) group size on the proportion of gargles and twargles directed towards the alpha  
 971 male. Monkey ID is a random effect.

	Age exp(est) *)	CI Age	SE Age	P-value	Group Size exp(est) *)	CI Grou p Size	SE Grou p Size	P-value
Gargle Model	1.094	1.06 7 - 1.21 2	0.01 3	<0.001	0.985	0.97 9 - 0.99 1	0.00 3	<0.001
Twargle Model	1.032	1.00 6 - 1.05 9	0.01 3	0.015	0.994	0.98 7 - 1.00 0	0.00 3	0.055

972

973 \*exp(est) is the exponential of the estimate reported by R's summary of the  
 974 glmmTMB function. In other contexts when only two groups are being compared we  
 975 refer to it as IRR. In this more general case, it is the ratio between the estimate of  
 976 the probability of the gargle being directed towards the alpha of n+1-year-olds vs.  
 977 n-yr-olds, or, in the case of group size, it is the ratio between the estimate of the  
 978 probability of the gargle being directed towards the alpha in group size n+1 vs. size  
 979 n. The confidence intervals refer to these quantities, i.e., 1.067 - 1.212 is the  
 980 confidence interval around 1.094.

981

982

983 For the gargle model, there are 1040 observations, 193 groups, and the variance in  
 984 the random effects intercept is 0.074. For the twargle model, there are 477  
 985 observations, 135 groups, and the random effects intercept variance is 0.010.

986

113

114

987 The raw data (Tables S8-9, below) show that for most ages and group sizes,  
 988 individuals gargle and twargle most frequently to the alpha male. Note that for age  
 989 5+, the proportion of gargles/twargles directed to the alpha male is well over 0.50  
 990 for all group sizes with a reasonable sample size. Data are scant for group sizes  
 991 below 10 and above 39.

992

993 Table S8: Proportions of gargles directed to the alpha male for different age groups  
 994 and group sizes (raw data). NA values indicate a complete absence of data for that  
 995 category. \* indicates a sample size of <10 gargles in that category, and † indicates  
 996 a sample of ≥100 gargles in that category. Age 0-1 indicates the first year of life.

997

**Proportion of Gargles directed to the Alpha Male by Group Size and Age**

998

Group Size

	<b>5-9</b>	<b>10-14</b>	<b>15-19</b>	<b>20-24</b>	<b>25-29</b>	<b>30-34</b>	<b>35-39</b>	<b>40-44</b>
<b>Age 0-1</b>	0.0*	0.69†	0.54†	0.63†	0.52†	0.47†	0.34†	NA
<b>1-2</b>	1.0*	0.40	0.82†	0.55†	0.39	0.51†	0.48†	0.36
<b>2-3</b>	1.0	0.74†	0.85†	0.91†	0.70	0.61†	0.59†	0.13*
<b>3-4</b>	NA	0.93	0.89†	0.83†	0.83	0.73	0.49†	NA
<b>4-5</b>	1.0*	1.0	0.92	0.97	0.77	0.11*	0.75	NA
<b>5+</b>	0.95	0.96†	0.88†	0.82†	0.85†	0.80†	0.81†	NA

1000

1001 Table S9: Proportions of twargles directed to the alpha male for different age groups  
 1002 and group sizes (raw data). NA values indicate a complete absence of data for that  
 1003 category. \* indicates a sample size of <10 twargles in that category, and † indicates  
 1004 a sample of ≥100 twargles in that category. Age 0-1 indicates the first year of life.

1005

**Proportion of Twargles Directed to the Alpha Male by Group Size and Age**

1006

Group Size

	<b>5-9</b>	<b>10-14</b>	<b>15-19</b>	<b>20-24</b>	<b>25-29</b>	<b>30-34</b>	<b>35-39</b>	<b>40-44</b>
<b>Age 0-1</b>	0.50*	0.75	0.66†	0.77†	0.66†	0.75	0.44	NA
<b>1-2</b>	NA	0.0*	0.71*	0.50	0.0*	0.73	0.92†	0.17*
<b>2-3</b>	1.0*	1.0	1.0*	0.89	0.67*	0.69	0.83	0.43*
<b>3-4</b>	NA	0.0*	0.78	0.82	0.98	0.83*	0.36	NA
<b>4-5</b>	NA	0.43*	0.93	0.93	0.92	0.0*	1.0*	NA
<b>5+</b>	0.92†	0.89	0.89†	0.87†	0.88†	0.82†	0.33*	NA

1008

1009 As group size increases to 30+ members, individuals gargle relatively less to the  
 1010 alpha male, especially when young (Table S8). We suspect this to be related to the  
 1011 threshold at which groups are prone to fission. In large groups, monkey “cliques”

115

116

1012 form, which may eventually separate into independent groups should a fission  
 1013 event occur. In the event of a fission, the current alpha male may no longer be a  
 1014 primary provider of valuable resources and protection, thus this vocalization  
 1015 dynamic may display bet-hedging behavior. However, the same trend is not true for  
 1016 twargles (Table S9). This suggests that gargles and twargles may serve distinct  
 1017 functions, although it is unclear what these distinctions are.

1018

1019 **Multiple Linear Regression Models with Pregnant Females and**  
 1020 **Females with Infants (<8 months) (Relevant to P2 and P3):** Pregnant  
 1021 females (P2) and females with infants (<8 months) (P3) were predicted to  
 1022 gargle and twargle at higher rates than the rest of the adult female  
 1023 population (>60 months and neither pregnant nor having an infant <8  
 1024 months old.)

1025 We ran multiple linear regression models with glmmTMB in R ([http://www.R-](http://www.R-project.org/)  
 1026 [project.org/](http://www.R-project.org/)). This approach differs from the modeling approach in the main  
 1027 text in that it includes the two reproductive states of interest - pregnancy  
 1028 and early lactation (i.e. having an infant <8 mo old) as fixed effects in the  
 1029 same model, rather than creating multiple separate models, each containing  
 1030 a single fixed effect. As in the other models, we include individual monkey ID  
 1031 as a random effect and include observation time as a log offset variable.

1032

1033 We predicted that pregnant females and adult females with infants (<8  
 1034 months) gargle and twargle more than all other adult (> 60 months)  
 1035 females. Our results show that pregnant females display slightly higher  
 1036 gargle rates, and females with infants (<8 months) display much higher  
 1037 gargle rates. (Gargle GLMM: n=306 observations, N=112 groups, random  
 1038 intercept variance=0.206; Twargle GLMM: n=297 observations, N=111  
 1039 groups, random intercept variance =0.611) (Table S10).

1040

1041

117

118

1042 Table S10: Results of multiple linear regression models in which (a)  
 1043 pregnancy and (b) having an infant <8 months old are fixed effects in the  
 1044 same model predicting gargle and twargle rates, in separate models.  
 1045 Incident rate ratios (IRR), exponentiated confidence intervals (CI), standard  
 1046 errors of the estimates (SE), P-values and Z-scores are presented for the  
 1047 fixed effects for GLMMs for Predictions 2 & 3.  
 1048

	IRR	CI	SE	P-value	Z-score	Prediction supported?
Gargle pregnancy variable in GLMM	1.486	1.087 - 2.032	0.160	0.013	2.484	weak
Gargle female with infant (<8 months) variable in GLMM	3.570	2.707 - 4.710	0.141	<0.0001	9.005	yes
Twargle pregnancy variable in GLMM	1.332	0.983 - 1.805	0.155	0.064	1.851	no
Twargle female with infant (<8 months) variable in GLMM	2.281	1.732 - 3.005	0.141	<0.0001	5.865	yes

1049

1050

```

119
120
1051 #This file contains the code used to run analyses in R, prepared with
1052 assistance from D. Cohen.
1053 #We explain the code, using the model presented in Table 3 as an
1054 annotated example for
1055 #interpreting the output.
1056
1057 $ R
1058 # ... omit output showing version, copyright, platform ...
1059
1060 > library(glmmTMB)
1061 # we use glmmTMB package
1062
1063 # Change the working directory, which must be set to where you (the user)
1064 saved the data files.
1065 > datadir=""
1066
1067 # Create the function to compute IRR and its confidence interval, which only
1068 needs to be defined once.
1069
1070 > expci <- function(coef, se){c(exp(coef),se*exp(coef),exp(coef +
1071 se*qnorm(.025)),exp(coef - se*qnorm(.025)))}
1072 # See description below of IRR computation.
1073
1074 # function (constructed after our data analysis) to show results of
1075 # R models constructed from imported data (again, only needs to be
1076 defined once)
1077 # More explanation appears after the first example.
1078
1079 > showresult <- function(model,datafile){
1080   data <- read.csv(paste(datadir,datafile,sep=""),na.strings="!@$%");
1081   m <-summary(glmmTMB(model, data=data,family=nbinom1));
1082   print(m);
1083   irr=expci(m$coefficients$cond[2,1],m$coefficients$cond[2,2])
1084   cat("IRR = ",irr[1]," CI: ",irr[3]," - ",irr[4],"\\n")}
1085
1086
1087 # Now an annotated first example:
1088
1089 > showresult(nvn~infant+offset(log(vnhr))+(1|focal),"P1_gargle.csv")
1090 Family: nbinom1 ( log )
1091 Formula:      nvn ~ infant + offset(log(vnhr)) + (1 | focal)
1092 Data: data
1093
1094     AIC     BIC logLik deviance df.resid

```

```

121
122
1095 2621.0 2637.0 -1306.5 2613.0 407
1096
1097 Random effects:
1098
1099 Conditional model:
1100 Groups Name      Variance Std.Dev.
1101 focal (Intercept) 0.7244  0.8511
1102 Number of obs: 411, groups: focal, 355
1103
1104 Overdispersion parameter for nbinom1 family (): 27.9
1105
1106 Conditional model:
1107      Estimate Std. Error z value Pr(>|z|)
1108 (Intercept) -0.89425  0.10291  -8.69  <2e-16 ***
1109 infant      2.34232  0.09795  23.91  <2e-16 ***
1110 ---
1111 Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
1112 IRR = 10.40539 CI: 8.587776 - 12.6077
1113
1114 # The text below explains how the output in the example above relates to
1115 # the
1116 # results presented in the manuscript.
1117 # showresult(nvn~infant+offset(log(vnhr))+(1|focal),"P1_gargle.csv")
1118 # the first argument, nvn~infant+offset(log(vnhr))+(1|focal)
1119 # is the model
1120 # each data row contains:
1121 # nvn - the number of gargles [would say nve for twargles]
1122 # infant - 1 if the row refers to an infant, and otherwise 0 [other models
1123 # have
1124 # different fixed effects, but the same regime follows,
1125 # i.e. a 1 if the group fulfills the criterion and a 0 if the group does not]
1126 # vnhrs - number of hours in which gargles could be observed by a trained
1127 # research assistant in the field [would say vehrs for twargle hours]
1128 # focal - id of monkey
1129 #
1130 # the second argument to showresult is the name of the data file
1131 #
1132 # In read.csv the na.strings argument is required in order to read
1133 # the monkey id NA as a regular ID rather than a missing id
1134 #
1135 # the function showresult
1136 # reads the data file
1137 # constructs the model (actually the summary of the model)
1138 # prints the summary

```

```

123
124
1139 # and finally computes the IRR and confidence interval with the
1140 # function expci.
1141 # The arguments of this function are the estimate and std.error
1142 # of the fixed effect, which are extracted from the summary.
1143 # (only the first fixed effect is shown - two of the models
1144 # below have two fixed effects, so require additional code.
1145 # Both models appear in the SI, not in the main text.)
1146
1147
1148 # We are mainly interested in the output line near the bottom:
1149 # infant    2.34232  0.09795  23.91  <2e-16 ***
1150 # showing the fixed effect of being an infant
1151 # (Other models have different fixed effects.)
1152 #
1153 # The paper shows the following results:
1154 # #observations and #groups are shown in the output above:
1155 # Number of obs: 411, groups: focal, 355
1156 # Z value is 23.91, in the z value column of the infant line
1157 # P value is shown as <2e-16, in the Pr(>|z|) column of the infant line
1158 # (the paper describes this as <.0001)
1159 #
1160 # In table 3 we also see:
1161 # SE, .098, corresponding to the Std. Error column of the infant line
1162 # (which says .09795, rounded to .098 in the table)
1163 # Var, .724 is shown in the Variance column of the line
1164 # focal (Intercept) 0.7244  0.8511
1165 # (again the value .7244 is rounded in the table to .724)
1166 # The other entries in table 3, IRR and CI, are computed by the
1167 # function expci with two arguments, the estimate and std. error of
1168 # the infant line.
1169 # IRR is exp (exponential function) of the estimate: e^(2.34232)=10.405.
1170 # CI is the last two values in the output of expci, showing the exponential
1171 # of the 95% confidence interval around the estimate, i.e., 95% confidence
1172 # interval of IRR.
1173
1174 # Below is the code for the remaining models, which should produce all of
1175 # the statistical results including IRR and CI
1176
1177 #P1 twargle
1178 > showresult(nve~infant+offset(log(vehr))+(1|focal),"P1_twargle.csv")
1179
1180 # Pregnant females will gargle and twargle at higher rates than the rest of
1181 # the adult female population, especially adult females who are not
1182 pregnant

```

```

125
126
1183 # and do not have an #infant <8 months old.
1184 #P2a gargle
1185 > showresult(nvn~pregnant+offset(log(vnhr))+(1|focal),"P2a_gargle.csv")
1186
1187 #P2a twargle
1188 > showresult(nve~pregnant+offset(log(vehr))+(1|focal),"P2a_twargle.csv")
1189
1190 #P2b gargle
1191 > showresult(nvn~pregnant+offset(log(vnhr))+(1|focal),"P2b_gargle.csv")
1192
1193 #P2b twargle
1194 > showresult(nve~pregnant+offset(log(vehr))+(1|focal),"P2b_twargle.csv")
1195
1196 #P3a gargle
1197 > showresult(nvn~nursing+offset(log(vnhr))+(1|focal),"P3a_gargle.csv")
1198
1199 #P3a twargle
1200 > showresult(nve~nursing+offset(log(vehr))+(1|focal),"P3a_twargle.csv")
1201
1202 #P3b gargle
1203 > showresult(nvn~nursing+offset(log(vnhr))+(1|focal),"P3b_gargle.csv")
1204
1205 #P3b twargle
1206 > showresult(nve~nursing+offset(log(vehr))+(1|focal),"P3b_twargle.csv")
1207
1208 # now we will use a different function for P4
1209 > showresult2 <- function(model,datafile,subset){
1210   data <- read.csv(paste(datadir,datafile,sep=""),na.strings="!@$%");
1211   m <-summary(glmTMB(model,
1212 data=eval(parse(text=paste("data[data$",subset,",]",sep=""))),family=nbino
1213 m1));
1214   print(m);
1215   irr=exp(m$coefficients$cond[2,1],m$coefficients$cond[2,2])
1216   cat("IRR = ",irr[1]," CI: ",irr[3]," - ",irr[4],"\n")}
1217
1218 #the difference between showresult2 and showresult is that there is an
1219 additional subset argument
1220 # which describes which subset of the data to use.
1221
1222 #P4 gargle
1223 > showresult2(nvn~fatherNOTalpha+offset(log(vnhr))+(1|
1224 focal),"P4_gargle.csv","X8mo==1")
1225
1226 # While the lines above, e.g., for P.1, compute a summary of a model like

```



```

127
128
1227 this:
1228 # > summary(glmmTMB(nvn~infant+offset(log(vnhr))+(1|focal),
1229 data=data,family=nbinom1))
1230 # showresult2 uses only a subset of the data, like this:
1231 # > summary(glmmTMB(nvn~fatherNOTalpha+offset(log(vnhr))+(1|focal),
1232 data=data[data$X8mo==1,],family=nbinom1))
1233 # In this case it is using only the data rows with column X8mo equal to 1
1234 # (in other words the data for monkeys less than 8 months old)
1235
1236 > showresult2(nvn~fatherNOTalpha+offset(log(vnhr))+(1|
1237 focal),"P4_gargle.csv","X24mo==1")
1238
1239 > showresult2(nvn~fatherNOTalpha+offset(log(vnhr))+(1|
1240 focal),"P4_gargle.csv","nursing==1")
1241
1242 > showresult2(nvn~fatherNOTalpha+offset(log(vnhr))+(1|
1243 focal),"P4_gargle.csv","pregnant==1")
1244
1245
1246 #P4 twargles
1247 > showresult2(nve~fatherNOTalpha+offset(log(vehr))+(1|
1248 focal),"P4_twargle.csv","X8mo==1")
1249
1250 > showresult2(nve~fatherNOTalpha+offset(log(vehr))+(1|
1251 focal),"P4_twargle.csv","X24mo==1")
1252
1253 > showresult2(nve~fatherNOTalpha+offset(log(vehr))+(1|
1254 focal),"P4_twargle.csv","nursing==1")
1255
1256 > showresult2(nve~fatherNOTalpha+offset(log(vehr))+(1|
1257 focal),"P4_twargle.csv","pregnant==1")
1258
1259
1260 #We needed a different function to look at the results for TableS10,
1261 # because we included more than one fixed effect.
1262 #This is almost the same as showresult except that it shows the IRR and CIs
1263 for
1264 # 2 separate fixed effects.
1265 # This model is relevant to both Predictions 2 & 3.
1266
1267 > showresult3 <- function(model,datafile){
1268   data <- read.csv(paste(datadir,datafile,sep=""),na.strings="!@#$$%");
1269   m <-summary(glmmTMB(model, data=data,family=nbinom1));
1270   print(m);

```

```

129
130
1271 irr=expci(m$coefficients$cond[2,1],m$coefficients$cond[2,2])
1272 cat(attributes(m$coefficients$cond)$dimnames[[1]][[2]],": IRR = ",irr[1],"
1273 CI: ",irr[3]," - ",irr[4],"\n")
1274 irr=expci(m$coefficients$cond[3,1],m$coefficients$cond[3,2])
1275 cat(attributes(m$coefficients$cond)$dimnames[[1]][[3]],": IRR = ",irr[1],"
1276 CI: ",irr[3]," - ",irr[4],"\n")
1277
1278 > showresult3(nvn~pregnant+nursing+offset(log(vnhr))+(1|
1279 focal),"Table_S10_gargles.csv")
1280
1281 > showresult3(nve~pregnant+nursing+offset(log(vehr))+(1|
1282 focal),"Table_S10_twargles.csv")
1283
1284 ### The groupsize model [Table S7] needs a slightly different function since
1285 it uses nbinom2
1286 > showresult4 <- function(model,datafile){
1287   data <- read.csv(paste(datadir,datafile,sep=""),na.strings="!@#$$%");
1288   m <-summary(glmTMB(model, data=data,family=nbinom2));
1289   print(m);
1290   irr=expci(m$coefficients$cond[2,1],m$coefficients$cond[2,2])
1291   cat(attributes(m$coefficients$cond)$dimnames[[1]][[2]],": IRR = ",irr[1],"
1292 CI: ",irr[3]," - ",irr[4],"\n")
1293   irr=expci(m$coefficients$cond[3,1],m$coefficients$cond[3,2])
1294   cat(attributes(m$coefficients$cond)$dimnames[[1]][[3]],": IRR = ",irr[1],"
1295 CI: ",irr[3]," - ",irr[4],"\n")
1296
1297 > showresult4(alpha ~ age5 + size + offset(log(total)) + (1 |
1298 id),"Table_S7_gargles.csv")
1299
1300 > showresult4(alpha ~ age5 + size + offset(log(total)) + (1 |
1301 id),"Table_S7_twargles.csv")
1302

```