

¹ Sex-dependent resource defense in a nectar-feeding bat

²
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⁸ **Abstract**

⁹ Aggressive defense can secure from the transient monopolization of a resource up to the long-term maintenance
¹⁰ of a territory. While such interference competition is common in nectar-feeding birds, reports in nectar-
¹¹ feeding bats are rare. *Glossophaga* bats have been observed to temporarily defend flowers but the extent
¹² of this monopolization, its effects on nectar intake, and underlying sexual differences remain unknown. We
¹³ investigated the resource defense behavior of 36 *Glossophaga mutica* in the laboratory. We presented bats
¹⁴ with two patches of computer-controlled artificial flowers and tracked individual nectar intake. Furthermore,
¹⁵ we established an automated method for detecting potentially aggressive interactions near the artificial
¹⁶ flowers. Theoretical models of interference competition predict more aggressive interactions when resources
¹⁷ are spatially more clumped. To test this, we varied resource distribution across two patches from clumped
¹⁸ to distributed and monitored bats' interactions in one male, one female, and four mixed-sex groups. Males
¹⁹ engaged in potentially aggressive interactions more often than females and in each group some individuals
²⁰ defended artificial flowers against others. Subordinate males experienced a substantial decrease in nectar
²¹ intake, while females were marginally affected by male aggression. These results suggest that aggressive
²² interactions and their effect on nectar intake are sex-dependent in *G. mutica*. Furthermore, aggressive
²³ interactions were more frequent and resource defense was only successful when resources were clumped. Our
²⁴ experimental set-up allowed us to perform an automated test of models of interference competition with a
²⁵ mammal under controlled laboratory conditions. This approach may pave the way for similar studies with
²⁶ other animals.

²⁷ **Lay summary**

²⁸ Males bully other males to get more food, but only when food is easy to defend. When flowers are spread
²⁹ out nectar-feeding bats rarely engage in fights. However, when there are rich flowers in one spot and no
³⁰ flowers elsewhere, some males start attacking others, denying them access to the nectar. Females do not seem
³¹ bothered by such male bullies, but when there are no males around, some females become bullies themselves.

³² **Keywords:**

³³ resource defense, economic defendability, bat, *Glossophaga*

³⁴ **1. Introduction**

³⁵ Competition for limited resources like food or mates is a ubiquitous phenomenon in the animal kingdom.
³⁶ Such competition can be indirect by exploiting a common resource and preventing others from benefiting

37 from it (Paton and Carpenter 1984); or it can be direct by aggressively defending a resource. The latter is
38 known as interference competition (Amarasekare 2002). Aggressive resource defense by excluding competitors
39 leads to priority of access to those resources and thus establishes dominance. One individual is dominant
40 over another if it directs aggressive behavior towards it (chasing, threatening, biting, etc.) while receiving
41 little or no aggression from the other (Chase et al. 2002). In the extreme, dominance behavior can lead to
42 exclusive territoriality. Territoriality is a concept belonging to an indivisible continuum starting with the
43 transient monopolization of a preferred feeding opportunity to the longer-term defense of an area as exclusive
44 territory. The rules of economic defendability (Brown 1964) determine the adaptive compromise to which
45 a species' dominance behavior will evolve and develop along this continuum. The establishment of feeding
46 territories is well known for nectar-feeding birds (Gill and Wolf 1975; Carpenter and Macmillen 1976; Boyden
47 1978; Ewald and Carpenter 1978).

48 The cost of defense, a key parameter in the economic defendability equation, is likely much higher for a
49 nocturnal, echolocating bat than for a diurnal, visually oriented bird. The successful resource defense is only
50 possible after the competition is detected. Visual detection in the daylight works well over long distances.
51 In contrast, for a nocturnal, echolocating bat, especially for phyllostomid bats that are able to echolocate
52 with whispering calls (Howell 1974; Hörmann et al. 2020; Yoh et al. 2020), detecting intruders at a feeding
53 territory's boundary would require expensive patrolling flights.

54 Within bats, the flower visitors have an additional advantage if they are intimately familiar with their
55 feeding area. Compared to an insect-hunting bat that must continually scan for elusive prey by active
56 echolocation, a flower visitor can approach a target with minimal echolocation when seeking specific flowers
57 at known locations (Thiele and Winter 2005; Winter and Stich 2005; Gonzalez-Terrazas et al. 2016; Rose et
58 al. 2016). Thus, it is not surprising that the longer-term defense of extensive feeding territories as commonly
59 observed in nectar-feeding birds is not known for glossophagine, nectar-feeding bats (but see Watzke 2006 for
60 nectar-feeding flying foxes). Nonetheless, several observations have documented aggressive food defense by
61 glossophagine bats. The inflorescences of *Agave desmettiana* with their copious nectar (Lemke 1985) may be
62 defended by males or females of *Glossophaga soricina* against conspecifics but only during some hours of the
63 night (Lemke 1984; Lemke 1985). When left unguarded, intruders quickly exploited the opportunity to feed
64 from the previously defended plants. The Costa Rican bat *Glossophaga commissarisi* occasionally defends
65 and temporarily monopolizes single inflorescences of the understory palm *Calyptrogyne ghiesbreghtiana* against
66 other hovering bats, perching bats and katydids (Tschapka 2003). A commonality in these reports was that
67 the defense did not cover the area of a typical feeding range but was restricted to a single or a few flowering
68 plants and was also limited to a small number of hours during the night. Thus, glossophagine bats can show
69 aggressive resource defense.

70 In this study, we investigated for a nocturnal, nectar-feeding mammal, the flower-visiting bat *Glossophaga*
71 *mutica*, the role of aggressive interactions for gaining access to nectar food. We tested the predictions of
72 models of resource defense (Grant et al. 2002) and interference competition (Grant 1993) during a naturalistic
73 foraging paradigm in the laboratory. The occurrence of resource defense is predicted to be highest at
74 intermediate levels of food abundance (Grant et al. 2002). In line with this prediction, the transient nature of
75 nightly defense behavior observed in the field suggests that changes in food-abundance or food-requirements
76 that occur within the night affected the strength of the observed behavior. To mimic the natural situation of
77 bat-pollinated flowers many of which replenish their nectar more or less continuously throughout a night (e.g.
78 Tschapka and Helversen 2007) we programmed artificial flowers to provide nectar with a fixed interval reward
79 schedule. Once a nectar reward had been taken by any bat, the fixed interval had to pass before the next
80 reward was available at this flower. Furthermore, theoretical models of interference competition predict that
81 clumped resources lead to more agonistic behavior and resource defense than evenly distributed resources
82 (Grant 1993). To include a test of this prediction in our experimental design, we spatially subdivided our
83 flower field into two patches and programmed them to automatically change during the night the spatial
84 distribution of available nectar resources. We performed our study with 36 individuals of male and female *G.*
85 *mutica*. By using artificial flowers in a closed environment, we could track all flower visits and total nectar
86 consumption of every individual in the group. Each individual carried an electronic ID tag and flowers were
87 equipped with ID sensors. This also enabled us to detect and quantify a typical class of aggressive interactions
88 between pairs of individuals directly at the artificial flowers fully automatically.

89 Our novel experimental set-up thus allowed us to perform a mostly automated experimental test of models
90 of interference competition and resource defense with a mammal under the controlled conditions of the

91 laboratory. This new approach may pave the way for further such studies with other groups of organisms.

92 **2. Materials and methods**

93 **(a) Subjects and housing**

94 Experiments were conducted with 36 individuals of the small, (9-10g) neotropical nectarivorous bat species
95 formerly identified as *Glossophaga soricina* (Pallas's long-tongued bat). In view of the recent taxonomic
96 revision of the *G. soricina* species complex (Calahorra-Oliart et al. 2021), it is relevant to note that the
97 founders of our colony used in this and all our previous studies were caught at the Cueva de las Vegas,
98 Municipio de Tenampulco, Mexico and transported to Germany in 1988. Thus they belong to the species *G.
99 mutica*. Bats came from our captive colony and were older than one year as judged by finger joint ossification
100 (Brunet-Rossini and Wilkinson 2009). They carried Radio Frequency Identification (RFID) tags attached to
101 cable tie collars (total weight of collar and RFID tag = 0.2g, max. 2.4% of the body weight) that were removed
102 after the experiment. Additionally, bats had numbered plastic split rings (A C Hughes Ltd., Middlesex, UK)
103 around the forearm for visual identification. Temperature in the experimental and colony room was kept at
104 20-25°C, air humidity at 65-75%, and light conditions were 12:12 LD (light off at 16h).

105 **(b) Experimental set-up**

106 In the experimental room ten artificial flowers with automated nectar delivery (Winter and Stich 2005) were
107 mounted along a 4.2m bar at a height of 1.2m (Fig. 1). The distance between flowers was 0.4m. Flowers were
108 divided into two groups of five to simulate two flower patches. Each patch was enclosed by a sheet-covered
109 frame around the four sides and at the top to separate the groups of flowers spatially (Fig. 1). The only
110 entrance to the patches was a 0.4m gap between the ground and the bottom end of this enclosure (Fig. 1,
111 dashed line). From this entrance bats had to fly up vertically to reach the flowers, which increased costs for
112 switching between patches. A stepper-motor syringe pump delivered nectar via tubes and pinch valves to
113 the artificial flowers. Nectar rewards were triggered by the interruption of an infrared light barrier at the
114 flower opening. The RFID reader below the flower head identified a bat's ID code. Flower visits (infrared
115 light barrier interruptions) and ID sensor events were recorded during every experimental night. The reward
116 schedule was configured using PhenoSoft Control (Phenosys GmbH, Berlin, Germany). Every detected event
117 at a flower (including date, time, individual ID, duration of the event and amount of nectar delivered) was
118 recorded for data analysis.

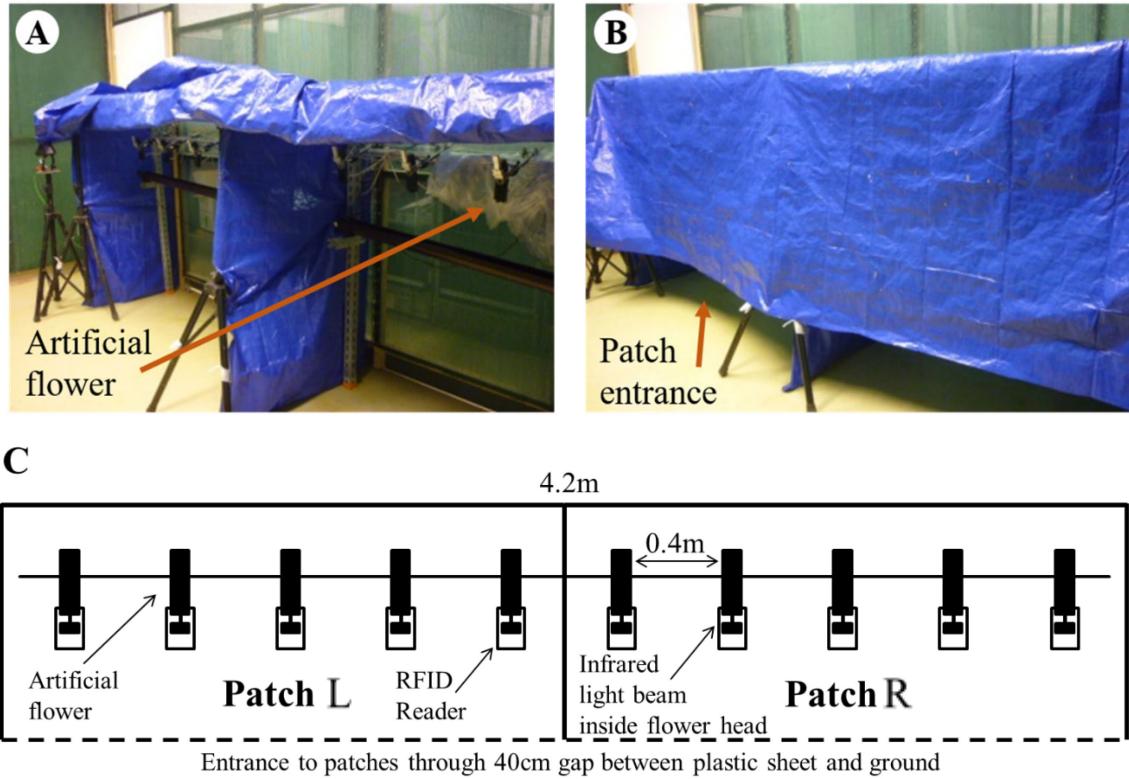


Figure 1: Experimental set-up consisting of two spatially separated patches of five flowers each. (A) The ten flowers were mounted 1.2m above ground. They were divided into two patches, L and R. (B) During experiments the patches were separated by plastic sheets. To make it more demanding for bats to enter a patch, the only entrance was through a 0.4m gap from the ground to the lower rim of the plastic foil. (C) Schematic drawing of the experimental set-up from above. The dashed line indicates the side with the patch entrance.

119 (c) Experimental procedure

120 Six bats were randomly caught from the colony and were tested simultaneously as a group. Four experimental
 121 groups consisted of three males together with three females (mixed groups), whereas one group consisted of
 122 six males, and another of six females. All bats were weighed before the experiment.

123 During the nightly experiments, in addition to the nectar provided by artificial flowers, bats had access to
 124 pollen and water and to 6mL of additional food containing 200mg NektarPlus (Nekton, Keltern, Germany)
 125 and 300mg milk powder (Milasan Folgemilch 2, Sunval Baby Food GmbH, Mannheim, Germany) dissolved in
 126 water. Rewards at flowers consisted always of 30 μ L nectar (15% w/w sugar concentration, sucrose: fructose
 127 1:2). Before the experimental schedule started, individuals were allowed to familiarize themselves with the
 128 set-up and the artificial flowers. Since during this training phase the plastic cover was removed, the two
 129 flower patches were not spatially separated and every flower visit was rewarded. This phase lasted for one to
 130 four nights until each bat visited the flowers regularly. One female of the first mixed group did not visit any
 131 artificial flower during the first night and was replaced by another female.

132 During the experiment, the two flower patches were covered and spatially separated (Fig. 1. Experimental
 133 nights were divided into two phases. During the first phase of the night only one of the two flower patches was
 134 rewarding, and therefore the resources were spatially clumped at a single location. The fixed time interval
 135 between rewards at each flower was 60s. During the second phase of the night both patches gave rewards,
 136 resources were evenly distributed across the two patches, and the fixed time interval between two rewards at
 137 a flower was increased to 120s. Therefore, the amount of food available per unit time did not change during
 138 the whole night; only the spatial distribution of food changed from the clumped resource condition with one

139 patch rewarding (five flowers) during the first phase of the night to the distributed resource condition with
140 two patches rewarding (ten flowers) during the second phase of the night. With this experimental schedule,
141 the maximal amount of nectar the bats could collect was $108mL$, which corresponds to $18mL$ nectar per
142 individual per night, roughly 150% of their daily requirement (Winter and Helversen 2001). The side of
143 the rewarding patch during the first phase of the night was chosen pseudo-randomly and the same patch
144 was never chosen in more than two consecutive nights. For the mixed groups, the duration of the clumped
145 resource condition was six hours and the experiment lasted nine nights (seven nights for the first mixed
146 group). For the same-sex groups, the duration of the first part of the night was variable (range = 4-8h, mean
147 = 6h) and the experiment lasted eight nights for the male group and seven nights for the female group.

148 (d) Chasing behavior

149 We took the frequency of individuals chasing each other at the artificial flowers as an indicator of the intensity
150 of aggressive interactions between group members. We developed a method to automatically detect and
151 score chasing events using the computer-collected animal identification data from the RFID sensors and
152 flower sensors. In a previous pilot study (Wintergerst 2018 Jan), three mixed groups of bats were video
153 recorded for 24h over 14 nights, and the video data were synchronized to the computer-collected data.
154 During this pilot study flowers were not covered by plastic sheets so that all flowers and the surrounding
155 room were visible on video. From the analysis of the combined data we were able to identify the following
156 pattern of visitation events that reliably indicated a chasing event between two identified individuals:
157 (i) an identified bat visited a flower and (ii) its visit was instantaneously followed by the detection of a
158 second bat, the chaser, that was detected very briefly (<200ms) and only by the ID sensor (detection
159 range 5-7cm). Importantly, this second bat never attempted to drink and therefore did not insert its nose
160 into the artificial flower and interrupt the light barrier inside the flower head. This distinguished such
161 a chase from the occasional quick succession of two feeding visits by two bats at the same flower. This
162 automated detection of chasing events not saves considerable time for the experimenter, but also avoids the
163 human observer bias, a common drawback in video analysis. For the 24 hours of combined video data and
164 automatically logged data, all 89 chasing events detected in the computer-logged data were confirmed by
165 video. Therefore, we consider the algorithm for detecting chasing events in the logged data to be highly
166 reliable. Of course, chases did not only occur at the artificial flowers. Thus, our chase numbers are only
167 an indicator of chasing intensity between pairs of bats. For example, in one hour of video we observed 61
168 chasing events, but only five of those occurred during flower visits and were also automatically detected.
169 However, since with our algorithm (see below) we detected a total of 1811 chasing events (35.5 ± 12.3
170 events per night during the experiment and only 4.8 ± 3 during the training nights, mean \pm SD) for
171 the 36 participating bats, we considered the automated approach adequate for quantifying within-group
172 dominance relationships. The total number of individual detections per night constrains the number of
173 chasing opportunities detectable with our method. Therefore, we corrected our counts of chasing events by
174 dividing the number of observed chases for each bat by the total number of detections for that bat on each night.

175

176 (e) Statistical analysis

177 To investigate the difference in chasing behavior between males and females and between the resource
178 conditions (one versus two rewarding patches) a Bayesian generalized linear mixed model (MCMCglmm, Hadfield
179 2010) with a binomial error structure was used. Body weight as an approximation of size and the full
180 interaction between resource condition and sex were included as fixed effects and the influence of these fixed
181 effects on the proportion of chasing events was assessed. Experimental group and individual were included
182 as random effects. The same model structure was used to address the question if the proportion of being
183 chased was influenced by these independent variables. If one or more individuals start to defend flowers and
184 thus exclude others from drinking, nectar consumption should increasingly differ between individuals since
185 the successful chaser should gain a higher nectar intake thus reducing the intake of the chased individuals.
186 Therefore, the between-individual difference in nectar consumption over the course of the experiment was
187 compared between experimental groups and resource conditions (clumped vs. distributed). First, each
188 individual's total nectar consumption standardized by the number of hours of foraging during the clumped

189 (one rewarding patch) and distributed (two rewarding patches) resource condition was determined for each
190 experimental night. Then these data were used to calculate group standard deviations, separately for the
191 males and females of each group. In order to assess the influence of resource defense on the individual
192 differences in nectar consumption (standard deviation of nectar intake) we fit a MCMCglmm model with a
193 Gaussian error structure and the following fixed effects: sex, experimental night (centered), and resource
194 condition (clumped or distributed), as well as all two-way interactions. Again, we included group and
195 individual as random effects.

196 By plotting individual nectar consumption during the last two nights of the experiment against the frequency
197 of chasing other individuals, two non-overlapping groups of males were obtained, which were labeled dominant
198 and subordinate males, respectively. Such a clear pattern was not observed in females. The identification
199 of dominant individuals was also supported by calculating the individual Glicko ratings (Glickman 1999;
200 So et al. 2015) from all chasing events over the last two experimental nights in each group. In the Glicko
201 Rating algorithm individuals gain or lose ranking points based on their wins or losses and the rating of their
202 opponent (Glickman 1999; So et al. 2015). Glicko ratings were analyzed using the PlayerRatings package
203 in R (Stephenson and Sonas 2020). Based on nectar consumption, the frequency of chasing events and the
204 individual Glicko group ranks (from 1 to 6, with 1 corresponding to the highest Glicko rating), each group
205 contained individuals belonging to one of three different types of social status: female, dominant male, and
206 subordinate male. To address the question whether nectar consumption varied depending on social status
207 during the early and late stages of the experiment we used Welch's tests and adjusted the p values using the
208 Holms method for multiple comparisons.

209 All statistical analyses were conducted using R (Team 2021). All data and code are available in the Zenodo
210 repository: xxx.

211 3. Results

212 (a) Example of nectar intake in one experimental group

213 The goal of our experiment was to investigate the sex-specific effects of resource defense in *Glossophaga*
214 *soricina*, in addition to the potential influence of interference competition on individual nectar intake. The
215 first striking observation we made was the uneven distribution of nectar consumed between the sexes and
216 individuals. For example, in the first mixed group of bats tested, after only two nights the nectar consumption
217 of two males was nearly reduced to zero, whereas the third male increased its consumption substantially (Fig.
218 2A). This pattern, however, only occurred during the clumped resource condition. Nectar consumption of
219 females did not change even during the clumped condition. On the same nights but during the second half of
220 the night, with resources distributed over two patches, nectar consumption of males and females converged at
221 the end of the experiment (Fig. 2B).

222 (b) Differences between sexes in frequency of chasing and being chased

223 In all mixed groups males chased other bats in front of flowers significantly more often than females did
224 (Fig. 3A, Table 1). Notably, the frequency of females as active chasers in female-only groups was higher than
225 chasing by females in the mixed groups (Fig. 3A). Although the rate of nectar availability remained constant
226 throughout the night and only the spatial distribution of the resources changed, the number of chasing events
227 was significantly lower during the distributed resource condition when rewards were available at both patches
228 (Table 1). There was no significant difference between the sexes in how often a bat was chased by another
229 individual (Fig. 3B) but individuals were chased less during the distributed resource condition (Table 1).
230 Weight as an indicator of size had no significant effect on chasing frequency or the frequency of being chased
231 (Table 1).

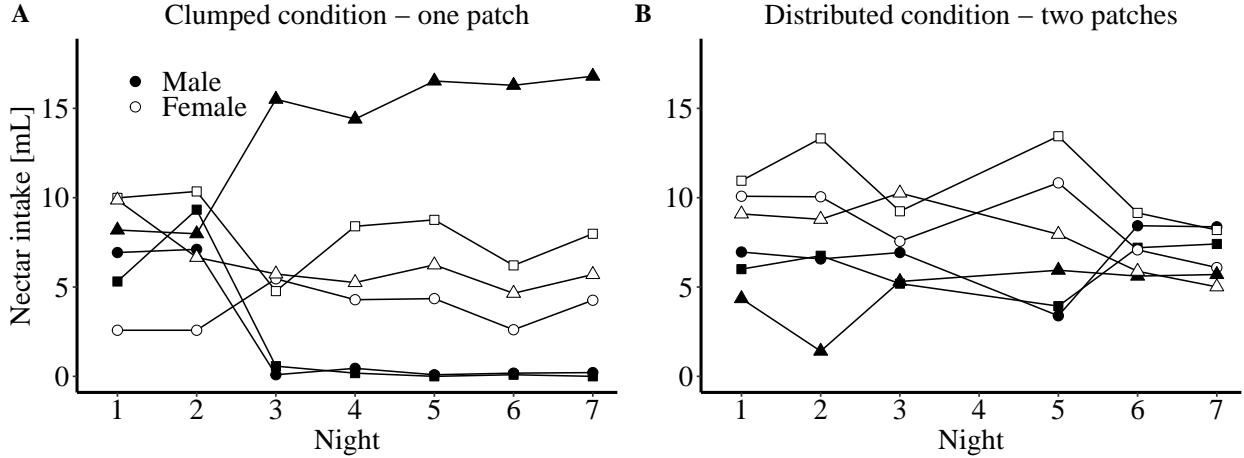


Figure 2: Change of individual nectar consumption from the clumped condition (A) to the distributed condition (B) during an experiment of one mixed group (3M, 3F, symbols show different individuals). (A). During the clumped resource condition (first part of the experimental night) rewards were only available at one patch. The nectar consumption of two subordinate males approached zero after only two nights, whereas the third, dominant, male greatly increased nectar intake during the experiment (males filled symbols). Females (open symbols) on the other hand maintained a stable level of nectar intake. (B) During the distributed resource condition (second part of the experimental night) rewards were available at both patches. Under this condition, individuals nearly equalized their level of nectar intake over the course of the experiment. The second part of night 4 was excluded due to technical problems.

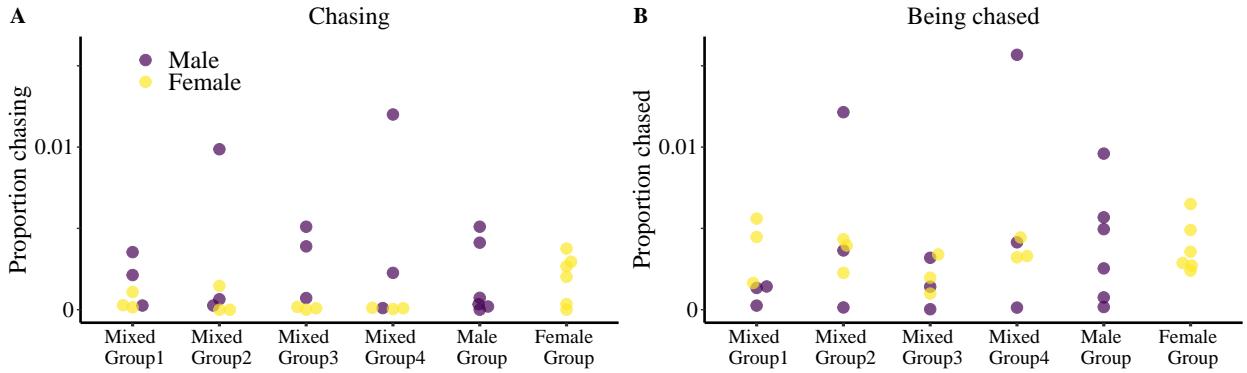


Figure 3: Sexes differed in the frequency of chasing or being chased during the clumped resource condition. (A) Males (dark symbols) chased others significantly more than females did (light symbols, Table 1). Shown are the individual proportions of chasing events over the whole experiment. Notably, in the females-only group some females chased more than any female in the mixed groups. (B) Being chased by other bats did not differ significantly between sexes (Table 1), but the variance of being chased was much higher for males.

Table 1: Summary of fixed effects from generalized linear mixed-effects models of chasing frequency and the frequency of being chased.

Model	term	estimate	95% credible interval	pMCMC
Chasing				
	(Intercept)	-6.52	(-18.06, 5.46)	0.252
	sex (female)	-2.09	(-3.47, -0.65)	0.001
	condition (distributed)	-0.49	(-0.76, -0.23)	0.001
	weight	-0.01	(-1.34, 1.1)	0.962
	sex (female):condition (distributed)	-0.29	(-0.75, 0.19)	0.228
Being chased				
	(Intercept)	-2.93	(-11.98, 4.61)	0.496
	sex (female)	0.90	(-0.12, 1.86)	0.078
	condition (distributed)	-0.98	(-1.24, -0.74)	0.001
	weight	-0.40	(-1.26, 0.44)	0.326
	sex (female):condition (distributed)	-0.13	(-0.48, 0.24)	0.440

Note: Fixed estimates whose credible intervals do not span zero are shown in bold. pMCMC = posterior probability

232 (c) Differences in nectar intake over time and between sexes and conditions

233 Resource defense should lead to a larger between-individual difference in nectar consumption (Brown 1964).
 234 Differences in nectar consumption were quantified as the standard deviation of nectar intake in each group,
 235 separately for males and females. During the clumped resource condition, the standard deviation increased
 236 significantly over time for males (Table 2, Fig. S1) and was generally higher for males than for females (Table
 237 2, Fig. S1). For females in the clumped resource condition the increase in standard deviation was significantly
 238 smaller than in males (significant interaction between sex and night, Table 2), and was not itself significant
 239 (estimate = 0.03, 95% CI = -0.02, 0.07). Compared to the clumped resource condition, in the distributed
 240 resource condition the effect of experimental night was significantly lower for males (interaction between
 241 condition and night, Table 2), but not for females (estimate = 0, 95% CI = -0.03, 0.02). Moreover, in the
 242 distributed resource condition there was no significant change over the course of the experiment in males
 243 (estimate = 0.01, 95% CI = -0.03, 0.05) nor in females (estimate = 0.03, 95% CI = -0.02, 0.06). Overall,
 244 for both males (significant effect of condition) and females (estimate = -0.07, 95% CI = -0.14, -0.02) the
 245 standard deviations were higher in the clumped than in the distributed resource conditions.

Table 2: Summary of fixed effects from a generalized linear mixed-effects model of the standard deviation of nectar intake over time.

term	estimate	95% credible interval	pMCMC
(Intercept)	0.52	(0.37, 0.68)	0.001
sex (female)	-0.20	(-0.26, -0.14)	0.001
condition (distributed)	-0.23	(-0.29, -0.17)	0.001
night	0.06	(0.01, 0.1)	0.022
sex (female):condition (distributed)	0.15	(0.06, 0.23)	0.001
sex (female):night	-0.03	(-0.06, -0.01)	0.015
condition (distributed):night	-0.04	(-0.07, -0.02)	0.001
sex (female):condition (distributed):night	0.04	(0.01, 0.08)	0.020

Note: Fixed estimates whose credible intervals do not span zero are shown in bold. pMCMC = posterior probability

246 (d) Social status and its effects on nectar intake

247 When plotting chasing events against nectar consumption the data for males fall into two non-overlapping
 248 groups. The males of one cluster (Fig. 4A, inside dashed oval) chased other individuals and consumed more
 249 nectar than the other males. This cluster included only one male from each of the four mixed groups but
 250 two males from the single males-only group. These six males were categorized as “dominant”. The second
 251 cluster of males (Fig. 4A, outside and below dashed oval) was characterized by a low frequency of chasing
 252 and low nectar consumption. These males were categorized as “subordinate”. In females such a pattern did
 253 not emerge (Fig. 4B). This classification was also supported by the Glicko ratings in each group (Fig. S2)
 254 and the observation that there was generally an inverse relationship between the frequency of chasing and the
 255 frequency of being chased, especially in males (Fig. S3). While in the females-only group four females chased
 256 other females more frequently, only one of these females would be classified as dominant using the same
 257 cut-off criteria we used for the males (Fig. 4B), but this was not the female with the highest Glicko rating
 258 (Fig. S2). During the last two nights of the experiment in the clumped resource condition, the highest nectar
 259 intake was observed in dominant males, with an intermediate intake in females, and lowest nectar intake
 260 in subordinate males (Fig. 5). In contrast, in the distributed resource condition there were no detectable
 261 differences between the nectar intake of dominant and subordinate males at any stage of the experiment (Fig.
 262 5), while the subordinate males had a significantly lower nectar intake than females in the first two, but not
 263 in the last two experimental nights (Fig. 5). Finally, the subordinate males increased their nectar intake from
 264 the clumped to the distributed condition, but the difference was only significant for the last two experimental
 265 nights (Fig. 5). While there was a correspondent decrease in the nectar intake of dominant males, it was not
 266 significant, most likely due to the small sample size ($n = 6$, Fig. 5). Again, in females there was no change in
 267 nectar intake between the resource conditions (Fig. 5)

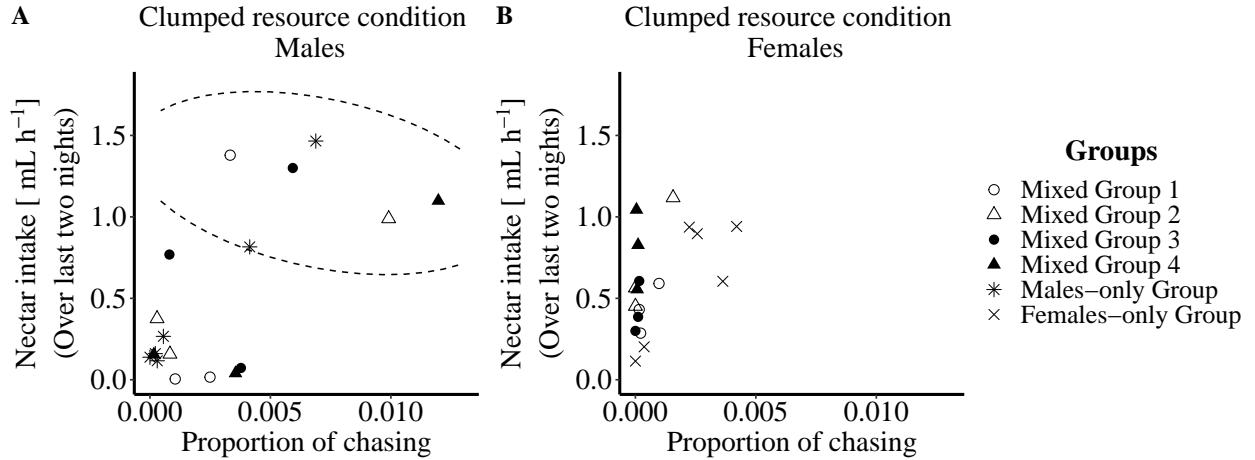


Figure 4: Influence of chasing frequency on nectar intake in the clumped resource condition during the last two nights of the experiment. **(A)** Males that more often chased other males also consumed more nectar. Males were divided into two non-overlapping groups by considering the chasing frequency and the amount of nectar an individual received during the clumped resource condition at the end of the experiment. Dominant males (inside dashed line oval) met two criteria: they chased other individuals at flowers more frequently (>0.003) and received more nectar ($>0.75\text{mL h}^{-1}$) during the clumped resource condition. Individuals outside the dashed line oval were categorized as subordinate males. **(B)** Nectar consumption of females did not generally depend on chasing frequency during the clumped resource condition and non-overlapping groups did not emerge.

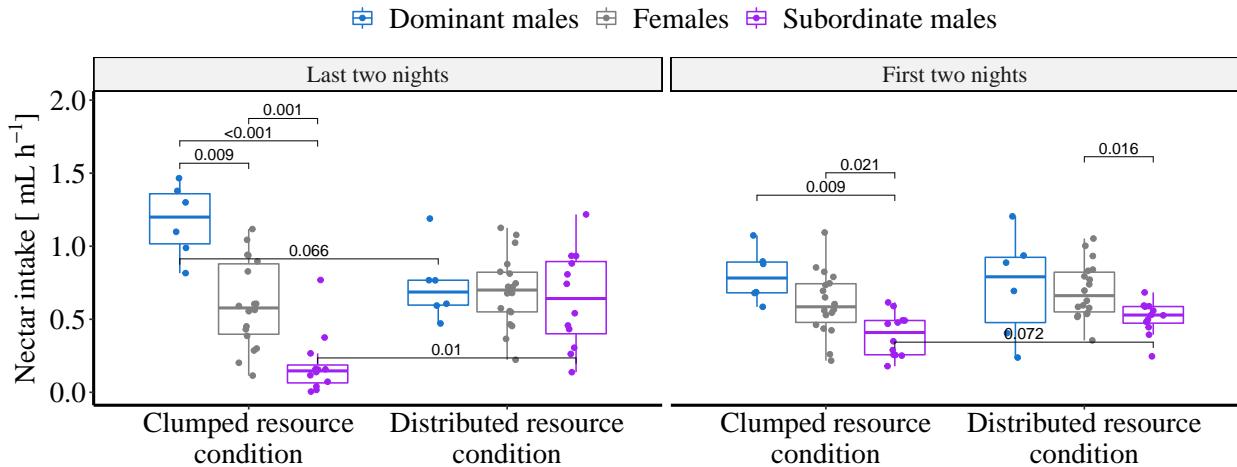


Figure 5: Comparison of nectar intake during the first and last two nights of the experiment depending on sex and social status. During the clumped resource distribution (left in each panel), already at the beginning of the experiment (right panel) subordinate males collected significantly less nectar than dominant males and females. At the end of the experiment (left panel), females, dominant and subordinate males differed to a large extent in their nectar consumption. During the distributed resource condition at the beginning of the experiment subordinate males received less nectar than females, but these differences disappeared by the end of the experiment. Numbers above brackets are the p values from unequal variance T tests (Welch's tests), adjusted for multiple comparisons using the Holms method. Contrasts between conditions were from paired Welch's tests. For clarity, only p values smaller than 0.1 are shown.

268 (e) behavioral observations

269 Qualitative behavioral observations of four hours of video recordings revealed several behaviors that seem to
 270 be characteristic for dominant males. Instead of just visiting the flowers and leaving the patch as the other
 271 individuals did, dominant males remained hanging between the flowers within the patch for a significant
 272 amount of time (Fig. S4). When other individuals came close due to visits of directly adjacent flowers,
 273 dominant males often spread one wing in the direction of the other individual which could be interpreted as
 274 a threatening posture. Some individuals were attacked and chased away by dominant males while visiting
 275 artificial flowers. In this case, dominant males mostly attacked from above with their mouth wide open, and
 276 followed the intruder for a short distance. Sometimes the chasing escalated into fighting with both bats
 277 tumbling towards the ground and resuming their flight only shortly above the floor. In rare cases, these fights
 278 might have led to small injuries. One subordinate male had several fresh scratches on its wing that were not
 279 present before the experiment and that were possibly caused by bites (Fig. S5). After a successful flower
 280 defense, the dominant male normally visited most of the five flowers within the patch before returning to its
 281 hanging position between the flowers.

282 4. Discussion

283 Similar to observations in free-living *Glossophaga* populations, in this experiment *G. mutica* competed for
 284 nectar not only by exploitation but also by interference competition. However, the results show that the
 285 predisposition to defend resources and the influence of interference competition on individual nectar intake
 286 differed significantly between the sexes. Only a subset of individuals, exclusively males in the mixed-sex groups,
 287 successfully defended flower patches. Dominant individuals were characterized by the highest frequency of
 288 chasing other individuals away from profitable flowers, by the highest Glicko ratings, and by a substantial
 289 increase in nectar intake during the time periods of active defense by the end of the experimental run.

290 Although the dominant males in the mixed groups chased females and other males equally often, only the
291 nectar intake of subordinate males but not of the females was affected by this behavior. Thus, male-initiated
292 interference competition increased the difference in nectar intake between males but did not affect females.
293 The frequency of aggressive interactions was higher, and males only defended resources successfully when
294 the available nectar was concentrated at only one flower patch. This supports the hypothesis that clumped
295 resources favor an increase in aggressive interactions (Grant 1993).

296 **(a) Sex-dependent resource defense and its differential effect on nectar intake,
297 depending on social status**

298 To our knowledge, this study is the first report of sex-dependent differences in resource defense behavior of
299 neotropical nectar-feeding bats. In mixed sex groups, females seemed to be much less affected by the behavior
300 of dominant males whereas subordinate males were excluded at least partially from the defended flower patch.
301 This finding is consistent with observations of free-flying *G. commissarisi*, in which males visited on average
302 a smaller number of artificial flowers than females did (Nachev and Winter 2019), presumably because of
303 interactions with other males. There are two possible explanations for this differential effect on subordinate
304 males and females. On the one hand, dominant males might just not be capable of excluding females. On
305 the other hand, dominant males could tolerate females in their defended patch because they might receive
306 additional benefits, for example tolerating females could lead to an increase in (future) mating opportunities.
307 Similar social dynamics have been described in the insectivorous bat species *Myotis daubentonii* (Senior et
308 al. 2005). Dominant males of this species temporarily exclude other males from profitable habitats whereas
309 females are tolerated and in addition to securing access to resources, the successful exclusion of other males
310 has been shown to increase the reproductive success of dominant males (Senior et al. 2005). Similarly, it has
311 been observed that male purple-throated carib hummingbirds (*Eulampis jugularis*), which successfully defend
312 highly profitable feeding-territories against other males while sharing the available resources with females,
313 experienced an increase in their mating success (Temeles and Kress 2010).
314 However, in our experiment dominant males chased females about as often as they chased subordinate males
315 (Table 1). If females were able to feed in the defended patch because dominant males tolerated them due
316 to potential additional benefits, it could be that the detected chasing events by dominant males differed in
317 quality depending on the sex of the intruder. This was not further quantified in the current study but could
318 potentially be investigated using audio recordings (Knörnschild et al. 2010). We extracted the frequency of
319 chasing events from data automatically recorded at artificial flowers (successive detection of two different
320 IDs while and after the first was feeding at the flower). Therefore, it was not possible to determine if males
321 showed behavioral differences when chasing other males in comparison to chasing females. The recorded
322 videos revealed that individuals chased each other not only directly at the artificial flowers but also in other
323 areas of the flower patch. Since individuals could only be identified by their ID tags directly at the ID reader
324 attached to artificial flowers the sex of individuals chasing each other in other areas of the experimental room
325 remained unknown. However, after the experiment some subordinate individuals showed marks from small
326 injuries at their wings (see example in Fig. S5) and such marks were only observed in males. This could be
327 an indication that dominant males directed more aggression (biting) towards subordinate males than towards
328 females. Such sexual dimorphism in aggressive resource defense is also known from other nectar-feeding
329 vertebrates, like hummingbirds. It is thought that the beaks of the males of some territorial hummingbirds
330 are specifically adapted as intrasexually selected weapons (Rico-Guevara et al. 2019).

331 **(b) Some observations from the single-sex groups**

332 Generally, females showed lower chasing frequencies, but, surprisingly, some females in the females-only group
333 showed an increased nectar consumption and chasing frequency, compared to the females in the mixed groups
334 (Fig. 4B). Thus it appears that in the absence of male individuals, some females exerted dominant behavior
335 over the other females, similar to males. These findings are similar to the social structure of resource defense
336 found in some nectar-feeding bird species. For example, in free-living ruby-throated hummingbirds females
337 also have lower levels of defense (Rousseau et al. 2014). Moreover, although both male and female *Eulampis*
338 *jugularis* hummingbirds defend feeding territories during the non-breeding season, males are always dominant
339 over females (Wolf and Hainsworth 1971; Temeles et al. 2005). It would be interesting to better understand

340 why females were less affected by the aggressive resource defense behavior of dominant males compared to
341 subordinate males and why females themselves did not consistently monopolize the profitable patch against
342 other females, not even in the females-only group. Of course, as there was but a single females-only group,
343 the latter observation needs to be confirmed with more data. One possibility is that females do not need to
344 defend flowers when a dominant male is already reducing the number of flower visitors and thus increasing
345 the amount of food available.

346 In all mixed sex groups, only one male per group became dominant and successfully defended flowers, whereas
347 in the males-only group two males exhibited dominant behavior (Fig. 4A). A closer look at the nectar
348 consumption at each flower revealed that on the last night of the experiment these two males had nearly
349 monopolized different flowers within the same patch rather than sharing access to the same flowers (Fig. S6).
350 Such flower partitioning was also observed in the females-only group (Fig. S7), but rarely seen in the mixed
351 groups (Figs. S8-S11). The successful resource defense by two individuals in the male-only group showed
352 that resource defense can occur independent of the presence of females, but, again, this was only based on a
353 single observation.

354 (c) Social status and social hierarchy

355 Although the position of the rewarding patch during the clumped resource condition changed between the
356 nights between the left and right, usually the same male continued to successfully defend the patch, especially
357 in the mixed-sex groups (Figs. S6-S11). This means that males defended the resources themselves and not a
358 particular location. Furthermore, this shows that even after changing the location of the defended patch
359 the same individuals were usually able to succeed in re-establishing their dominance against other males,
360 indicating a stable hierarchy at least for the duration of the experiment.

361 The ability of an individual to successfully defend and monopolize resources is often correlated with distinct
362 physical characteristics for example body size (Searcy 1979). However, in our results weight as an approxima-
363 tion of size did not correlate significantly with the chasing frequency of individuals (Table 1) and therefore
364 did not predict which male succeeded to defend a flower patch. Another factor that could influence the
365 success in defending flowers is age and therefore experience (Yasukawa 1979; Arcese 1987). Since we could
366 only discriminate between young and adult animals, we cannot dismiss age and experience as a predictor of
367 successful flower defense.

368 In this study, subordinate males received considerably less nectar than dominant males and females (Fig. 5).
369 However, except in mixed group 1, subordinate males were rarely completely excluded from the flower patch
370 and their average nectar intake during the clumped resource condition was still $0.3 \pm 0.18 \text{ mL h}^{-1}$ (mean \pm
371 SD). This result is in accordance with observations of free-living *G. soricina* in Colombia. There, subordinate
372 bats exploited the flowers defended by other individuals as soon as the dominant bat temporarily ceased
373 defending (Lemke 1984). Furthermore, in our study the frequency of chasing events decreased significantly
374 during the distributed resource condition in the second part of the night (Table 1). This supports the
375 theoretical prediction that aggressive defense behavior increases when resources are spatially concentrated
376 (Grant and Guha 1993), with the important caveat that the sequence of conditions was not controlled in this
377 experiment. With the current data we cannot answer whether the dominant males would successfully defend
378 a patch if the condition changed from distributed to clumped, but we believe this is a different question that
379 should be addressed separately. Resource defense should only occur when the energy gain outweighs the
380 cost of aggressive interactions (Brown 1964). Thus, our results could be explained by the decrease in quality
381 of the defended patch once its nectar supply rate dropped to half. This is also supported by the very low
382 number of chases observed during training when the flowers gave unrestricted rewards and were not separated
383 in discrete patches. Together, these results suggest that along the different degrees of territorial behavior,
384 resource defense observed in *Glossophaga* seems to represent a transient monopolization of resources instead
385 of a longer-term permanent exclusion of intruders.

386 (d) Conclusion

387 Although flower defense behavior of *G. mutica* was investigated in a laboratory setting, we observed similar
388 behavior as described in free-living *Glossophaga* populations. Our results revealed a sexual dimorphism in
389 flower defense behavior in mixed-sex groups. Only males successfully defended flower patches and excluded

390 other males from their defended resource, whereas females remained unaffected by this male behavior and
391 continued to visit the flowers guarded by a male. This observed pattern is similar to resource defense behavior
392 observed in other nectar-feeding vertebrates. Furthermore, we could show that the frequency of aggressive
393 interactions was, as predicted, higher when resources were clumped in one patch and transient. Future studies
394 with free-living populations have to be conducted to assess how frequent and important resource defense in
395 these nectar-feeding bats is and if males that are successful in defending resources have additional fitness
396 advantages.

397 **Supplementary material**

398 **Video analysis**

399 There were 89 chase occurrences observed ($f \rightarrow f$ 4 times, $f \rightarrow m$ 2 times, $m \rightarrow f$ 59 times, $m \rightarrow m$ 24 times).
400 Every time the algorithm marked an event as a chase event, there were two individuals following each other.
401 Some chase sequences did not get detected. The individual that chased never drank immediately after the
402 chase at the same flower location where the chase occurred. There were 16 incidences that were difficult to
403 classify by observation or did not appear to be aggressive interactions.

404

405 $f \rightarrow f$ appear to be less aggressive

406 $f \rightarrow m$ appear aggressive

407 $m \rightarrow f$ appear aggressive

408 $m \rightarrow m$ appear aggressive

409

410 **Supplementary figures**

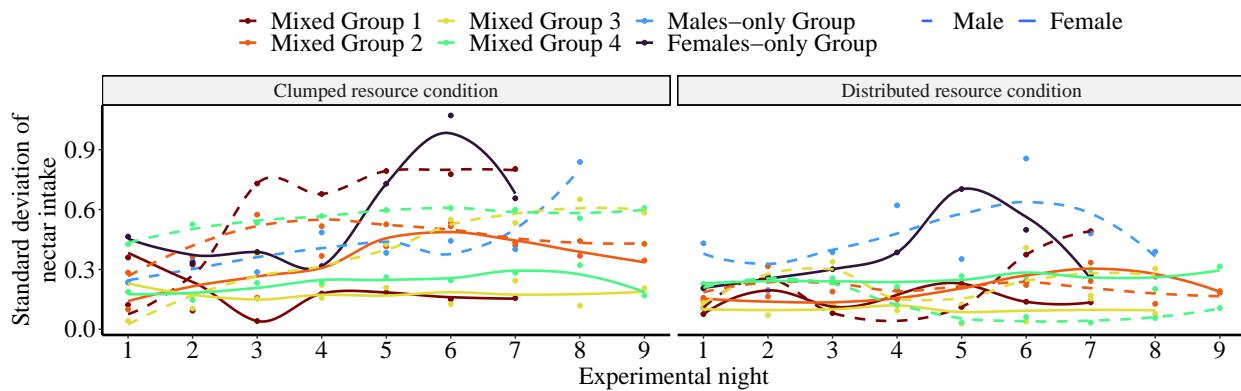


Figure S1: The standard deviation of group nectar consumption was used to measure the between individual differences in nectar intake. It was calculated for the clumped (left panel) and the distributed (right panel) resource conditions, separately for males (dashed lines) and females (continuous lines) from each experimental group (different colors). For visualization only, lines give the corresponding fits based on locally weighted scatterplot smoothing (loess). The statistical analysis was based on linear regression (see Methods).

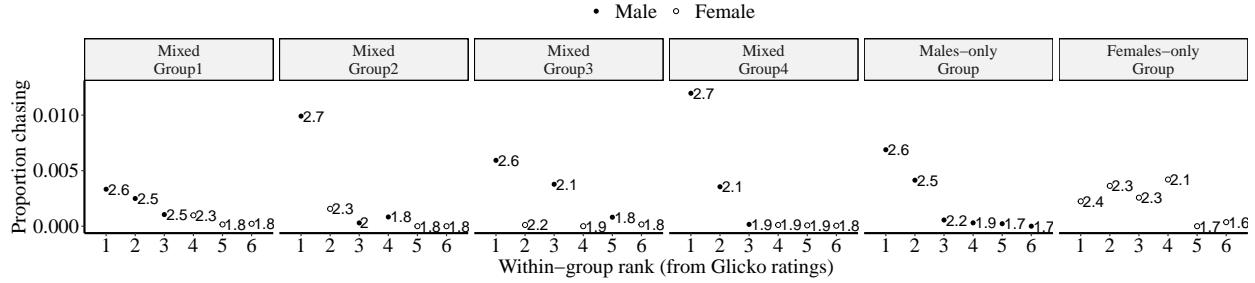


Figure S2: Glicko ratings within the six experimental groups. Over the last two experimental nights, the males (closed symbols) with the highest proportion of chasing events were also the individuals with the highest Glicko rating in each group (panels) during the clumped resource condition. There was no such correspondence for females in the female-only group (right panel). Numbers at symbols give the Glicko rating in thousands.

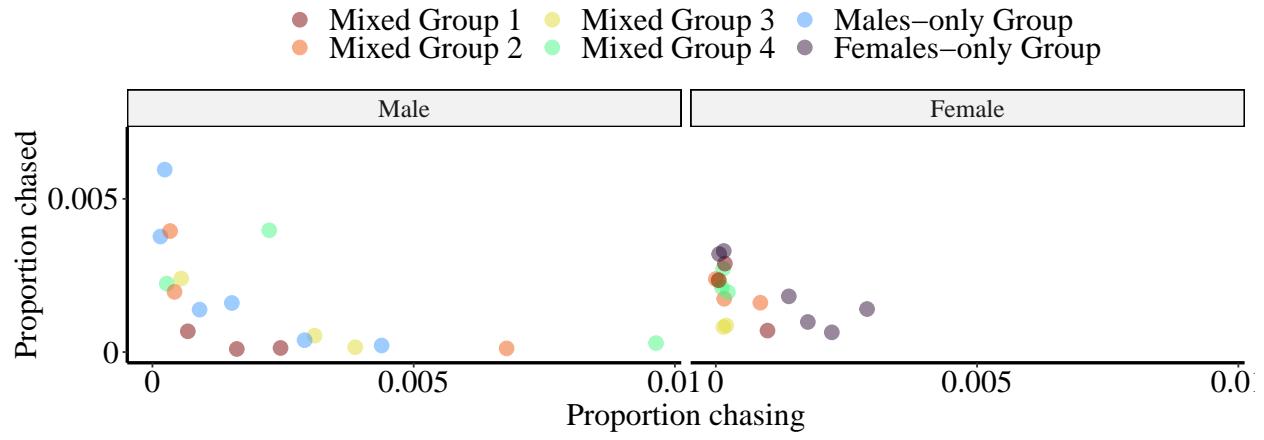


Figure S3: The relationship between proportion of chases versus the proportion of being chased out of all detections for female (right) and male (left) individuals in the six experimental groups (different colors).

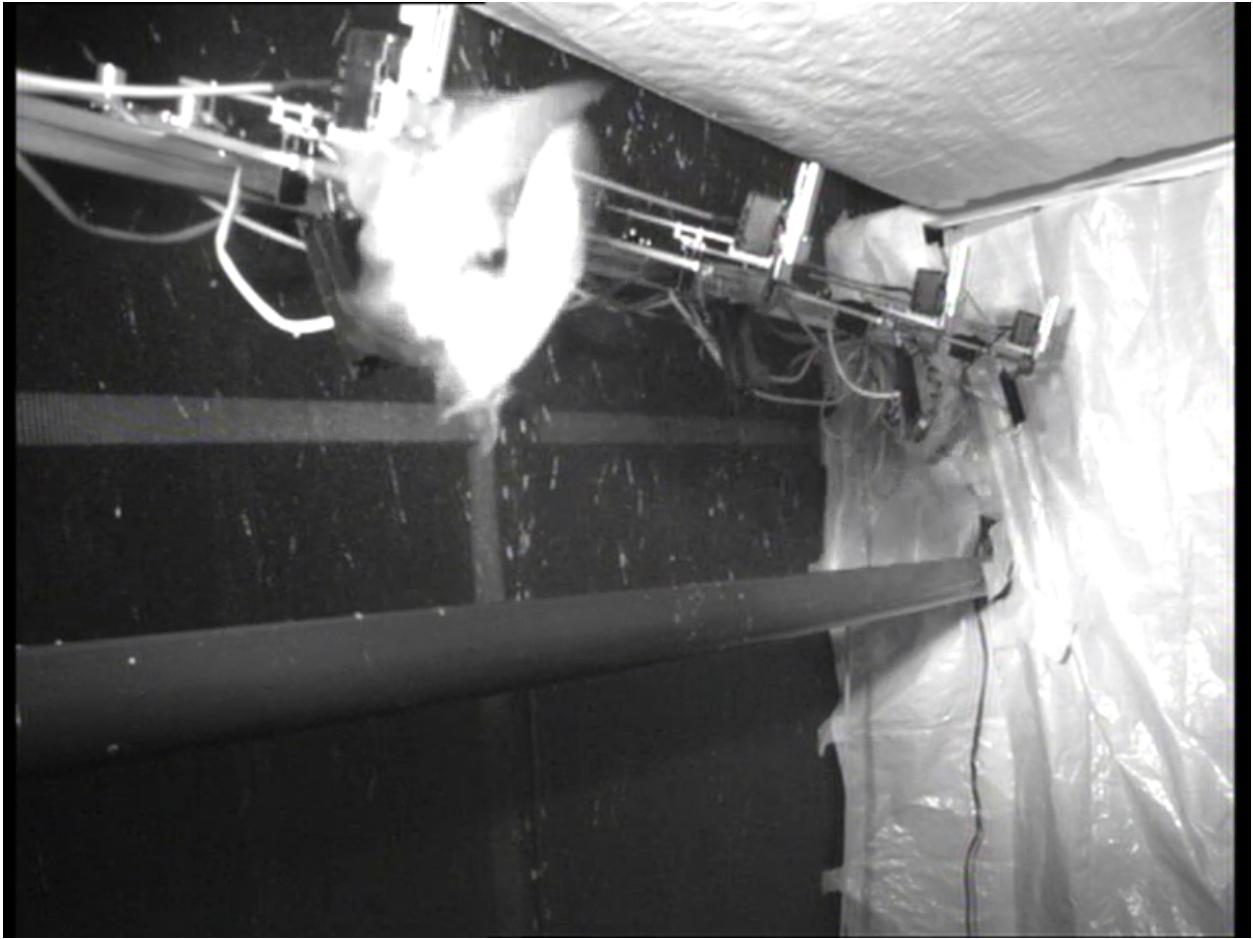


Figure S4: Video of the dominant male in mixed group 3 chasing all bats approaching the rewarding flowers in the rewarding patch during the clumped resource condition.

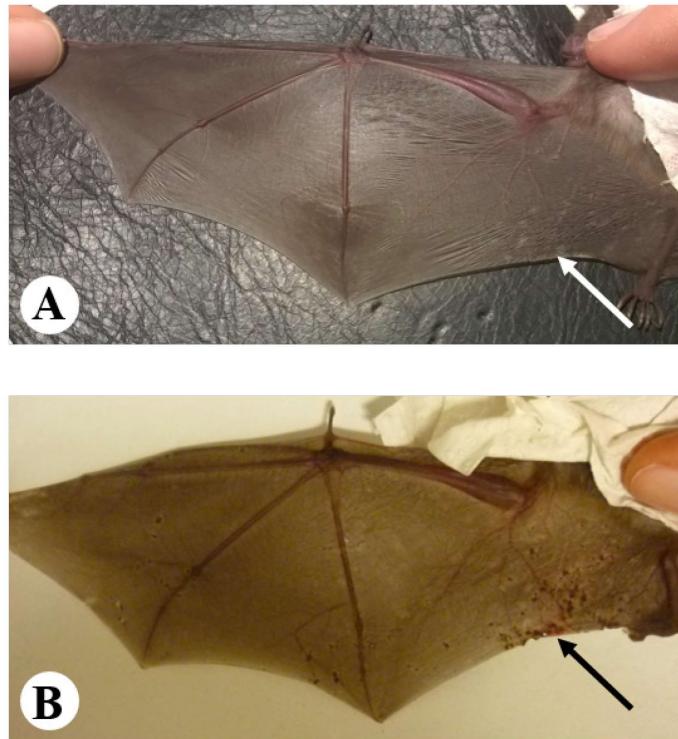


Figure S5: Wing images of a subordinate male from mixed group 4. The same individual was photographed before (**A**) and after the experiment (**B**). The black arrow points to the scarred location due to wing injuries, purportedly caused by the dominant male.

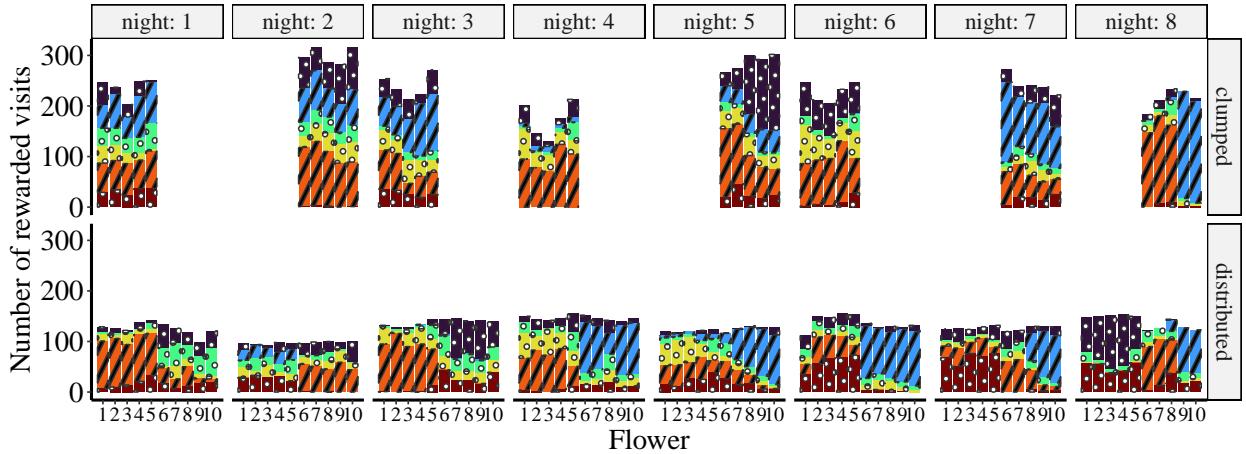


Figure S6: Distribution of rewarded visits across flowers for the six bats in the males-only group. The colored bars give the number of rewarded visits of each individual at the ten flowers during the clumped (top) and distributed (bottom) resource conditions for each experimental night (columns). The dominant males are shown with black stripes and the subordinate males are shown with white dots. This was the only group with two males behaving as dominant. On the last night, rather than sharing all flowers within the defended patch, the dominant males partitioned the patch into two subpatches, with each bat defending its own partition.

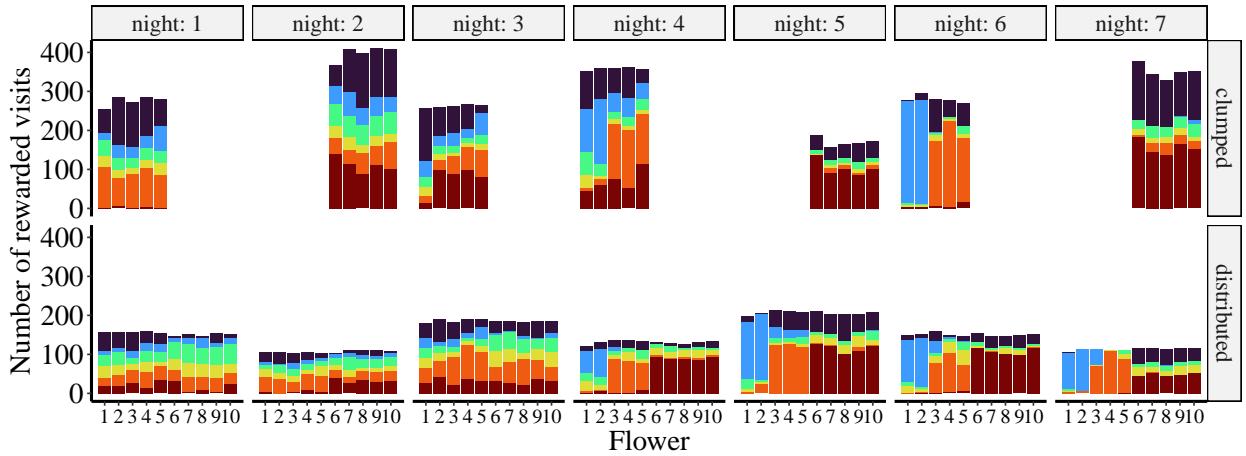


Figure S7: Distribution of rewarded visits across flowers for the six bats in the females-only group. The colored bars give the number of rewarded visits of each individual at the ten flowers during the clumped (top) and distributed (bottom) resource conditions for each experimental night (columns). Females in this group exhibited the highest frequency of chasing behavior compared to all other females. This is also the only group, in which females nearly monopolized flower patches or flowers within a patch.

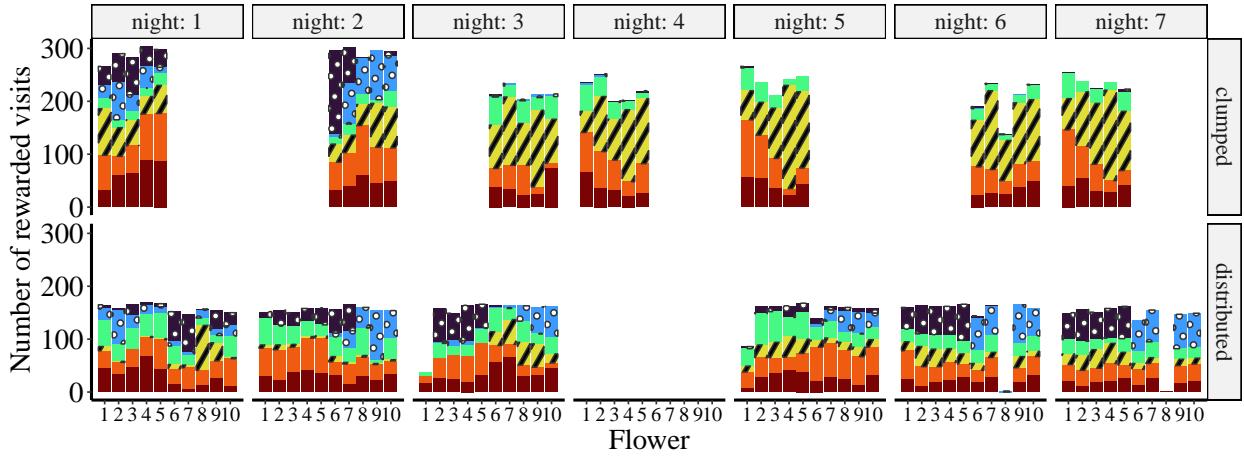


Figure S8: Distribution of rewarded visits across flowers for the six bats in mixed group 1. The colored bars give the number of rewarded visits of each individual at the ten flowers during the clumped (top) and distributed (bottom) resource conditions for each experimental night (columns). The dominant male is shown with black stripes, the subordinate males are shown with white dots, and the females are shown with solid bars. Due to a technical malfunction on night 4, there were no rewards delivered in the distributed resource condition and the data were excluded from analysis.

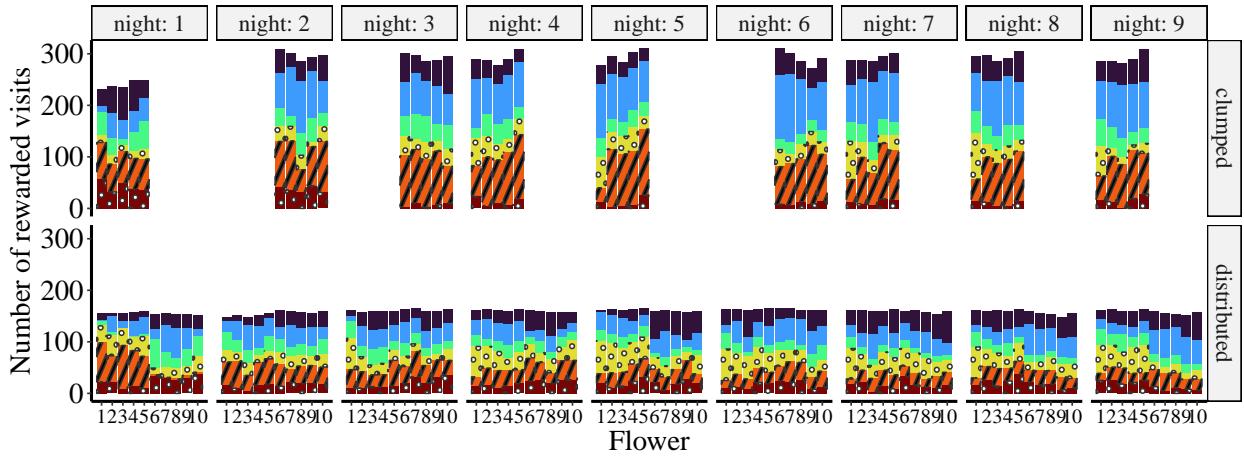


Figure S9: Distribution of rewarded visits across flowers for the six bats in mixed group 2. Same notation as in Fig. S8, but the colors correspond to different individuals.

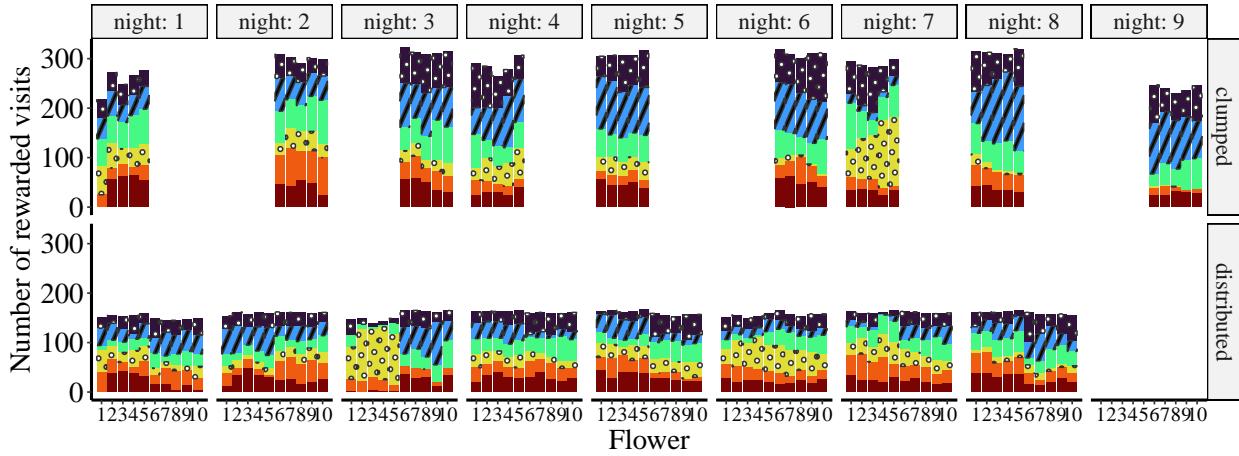


Figure S10: Distribution of rewarded visits across flowers for the six bats in mixed group 3. Same notation as in Fig. S8, but the colors correspond to different individuals. Due to a technical malfunction on night 9, there were no rewards delivered in the distributed resource condition and the data were excluded from analysis.

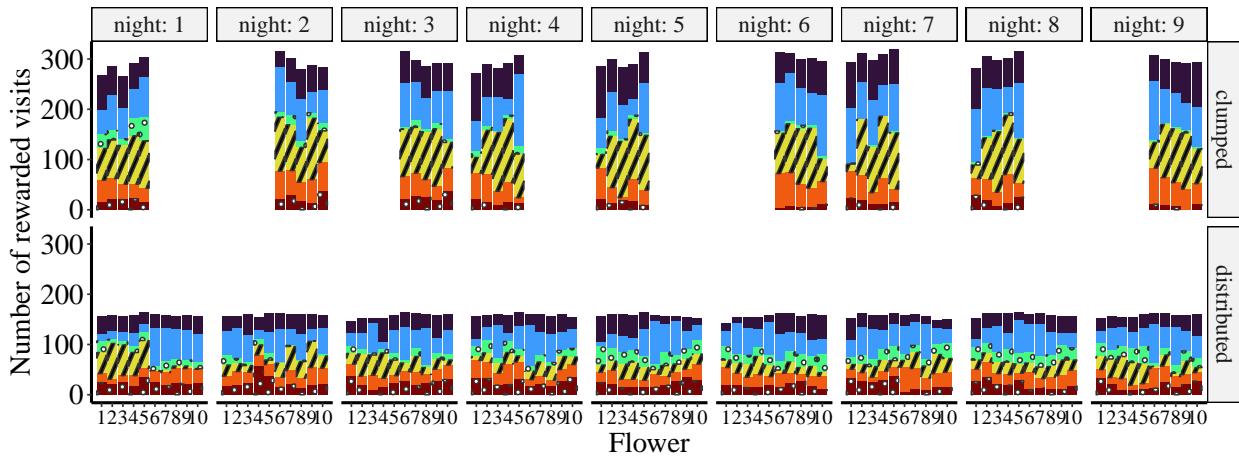


Figure S11: Distribution of rewarded visits across flowers for the six bats in mixed group 4. Same notation as in Fig. S8, but the colors correspond to different individuals.

411 Authors' contributions

- 412 412 S.W. Conceptualization, Methodology, Software, Data collection, Formal Analysis, Video Analysis, Writing—
original draft. V.N. Conceptualization, Methodology, Software, Formal Analysis, Data curation, Writing—
review and editing, Visualization, Supervision, Project Administration.
- 413 413 Y.W. Conceptualization, Resources, Methodology, Software (data acquisition), Writing—review and editing,
Supervision, Funding.

417 Competing interests

- 418 418 We declare we have no competing interests.

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426 **Data Availability**

427 All data and code are available in the Zenodo repository: xxx.

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