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Diet composition of the gecko species, *Cyrtodactylus leegrimeri*, on Hon Khoai Island, southern Vietnam

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17 **Diet composition of the gecko species, *Cyrtodactylus leegrimeri*, on Hon Khoai Island,**
18 **southern Vietnam**

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35

36 **Abstract.** The study investigated the diet composition of *Cyrtodactylus leegrismeri* on Hon Khoai
37 Island, southern Vietnam. A total of 852 prey items representing 25 prey categories were identified
38 from the stomach contents of 128 individuals. The diet of *C. leegrismer* included small
39 invertebrates, with Isoptera (termites) being the primary prey category, followed by Orthoptera,
40 Blattodea, Lepidoptera, Araneae and other prey groups. The species is characterized with a high
41 degree of dietary specialization, consuming a narrow spectrum of prey. There are no significant
42 differences in the overall diet composition of *C. leegrismeri* between seasons (rainy vs. dry) and
43 between habitats (forest vs. road). However, the results documented a lower prey richness in
44 disturbed habitats, with an increased reliance on termites, compared to the prey in forest habitats.
45 Regarding conservation, we highlight the importance of forest protection for safeguarding fauna
46 diversity on Hon Khoai Island.

47 **Keywords.** Dietary specialization, Disturbed habitats, Forest, Important index, Seasons, Termites.

49 Understanding dietary composition is essential in ecological research, as it reveals how
50 organisms adapt to their natural habitats (Lihoreau et al., 2015). As a key element of life history,
51 the diet shapes patterns of energy allocation, survival, reproduction and long-term evolutionary
52 adaptation (Sibly et al., 2012; Barnagaud et al., 2019). Thus, the spatial variation in diet
53 composition can be particularly informative in explaining the local adaptation of an endemic species
54 in restricted areas, including islands (Heaney, 2007; Whittaker et al., 2008; Lomolino et al., 2010;
55 Ngo et al., 2024).

56 With a high level of endemism, the third-largest vertebrate genus – *Cyrtodactylus*, comprising
57 nearly 400 recognized species worldwide, represents an exceptionally complex taxonomic group
58 (Grismer et al. 2021b; Uetz et al., 2026). The diversification of *Cyrtodactylus* species can be better
59 understood through integrative taxonomic approaches, in combination with ecological niche
60 studies, including dietary niche research (Sibly et al., 2012; Barnagaud et al., 2019; Grismer et al.,
61 2021a, 2021b; Ngo et al., 2022; 2025). *Cyrtodactylus leegrimeri* Chan and Norhayati, 2010, was
62 first discovered from a small Malaysian island and reported afterwards from several islands of
63 southern Vietnam, including Hon Khoai Island, Ca Mau Province (Chan and Norhayati, 2010;
64 Grismer and Grismer, 2017; Ngo et al., 2022). The oceanic barrier together with adaptation of
65 populations to distinct niches, including diet composition, might shape the distribution of *C.*
66 *leegrimeri* on the islands (Hutchinson, 1957; Heaney, 2007; Soberón, 2007). Recently, surveys on
67 Hon Khoai Island by Ngo et al. (2024) documented the diet composition of *Cnemaspis*
68 *psychedelica*, occurring in syntopy with *Cyrtodactylus leegrimeri*.

69 The natural forest on Hon Khoai Island has been recently degraded by human activities,
70 leading to a decline in reptile populations (Ngo et al., 2016b, 2018, 2022). The species of *C.*

71 *leegrimeri* and other reptiles were even soaked together in rice wine and used for traditional
72 medicine by road workers on Hon Khoai Island (Ngo et al., 2018). Such impacts are thought to
73 further influence both their populations and abundance of prey on the island (Ngo et al., 2024). In
74 the present study, we conducted field surveys on the island, and collected stomach contents of *C.*
75 *leegrimeri* in both forest and road habitats. Prey items were identified and compared between the
76 rainy season (November 2015) and the dry season (January 2016 and March 2025), and between
77 forest and disturbed habitats. Based on the findings, we examined whether seasonal conditions and
78 human disturbance impact on the species in terms of food selection.

79

80

MATERIAL AND METHODS

81 *Study site*

82 Hon Khoai Island is located in Rach Gia Bay, approximately 14.6 km from the mainland of
83 Ca Mau Province in southern Vietnam. It is the largest of 92 islands in the Bay, with the highest
84 peak of 318 m above sea level and a total area of approximately 4 km² (Ngo et al. 2022). The
85 terrestrial ecosystem is primarily characterized by granitic formation and dense forest with small
86 to medium-sized trees (Ngo et al., 2018, 2022). Ngo et al. (2022) recorded a checklist of 19
87 herpetofaunal species on Hon Khoai Island, including the focal species of *Cyrtodactylus*
88 *leegrimeri*. The species was observed nocturnally across a range of microhabitats, including
89 granite boulders, branches, leaves on vegetation 0.1–2 m above ground, and the forest floor (Ngo
90 et al. 2022).

91

92

93 *Field surveys*

94 Field surveys were conducted in forest habitats and along road trails on Hon Khoai Island,
95 Ca Mau Province, southern Vietnam, during the rainy season in November 2015 (forest), and the
96 dry season in January 2016 (forest) and March 2025 (road) (Fig. 1). Nocturnal excursions were
97 carried out after sunset between 19:00 and 24:00. Individuals were captured by hand and released
98 at the same spot after morphometric measurements and obtaining stomach contents. We identified
99 *C. leegrismar*, based on the following combination of characters: maximum snout-vent length
100 (SVL) up to 92 mm; conical, keeled tubercles on the occiput, body, hind limbs, and tail base (absent
101 on forelimbs); 18–19 longitudinal tubercle rows and 27–35 ventrals at midbody; a single row of
102 enlarged median subcaudals; transversely expanded subdigital lamellae (18–20 under fourth toe);
103 four precloacal pores in an arch, separated from a single row of enlarged, poreless femoral scales;
104 absence of a precloacal groove but presence of a depression; and a dorsal pattern of brown
105 coloration with a dark-brown stripe forming a “V” on the nape, dark head reticulations, irregular
106 dark-brown bands on body and tail, and mottled dark-brown limbs (Chan and Norhayati, 2010;
107 Ngo et al. 2022).

108

109 *Stomach content analysis*

110 A stomach-flushing technique was used to obtain stomach contents without sacrificing
111 animals (Griffiths, 1986; Norval et al., 2012; Nguyen et al., 2018; Ngo et al. 2024; 2025). Prey
112 items were collected by stomach-flushing with fresh water, using spatula, forceps, two syringes
113 with thread (60 ml), and an infusion tube of soft material (silicon). All collected items were
114 preserved in 70% ethanol and deposited in a refrigerator at a temperature of -4°C at the Institute
115 of Biology (IB), Hanoi, and Hong Duc University, Thanh Hoa Province, Vietnam.

116 In the laboratory, prey items were identified to the lowest possible taxonomic level, using a
117 stereomicroscope, following CSIRO (1991), Thai (2003) and Triplehorn et al. (2005). The
118 maximum length (L) and width (W) of each prey item were measured to the nearest 0.01 mm using
119 either a digital caliper for large prey or the stereomicroscope with a camera for small prey (Hirai
120 and Matsui, 2001). The prey volume (V, mm³) was calculated using the formula for a prolate
121 spheroid ($\pi = 3.14$; Magnusson et al., 2003): $V = 4\pi/3 \times (L/2) \times (W/2)^2$. The index of relative
122 importance (Ix), was used to determine the importance of each food category (except for
123 unidentified categories) and was calculated following the formula: $Ix = (\%F + \%N +$
124 $\%V)/3$ (Caldart et al., 2012), where $\%F = (S_i/S_t) \times 100$ (F : frequency of occurrence; S_i :
125 number of stomachs with the prey category; S_t : total number non-empty stomachs), $\%N =$
126 $(N_i/N_t) \times 100$ (N : number; N_i : number of items of a prey category; N_t : total number of of all prey
127 items); $\%V = (V_i/V_t) \times 100$ (V_i : volume per prey category per stomach; V_t : total volume of prey
128 in all stomachs).

129 Furthermore, we applied the “Dominance” index: $D = \sum[n_i(n_i - 1)]/[N(N - 1)]$, where n_i
130 is the number of food items in the i th taxon category and N is the total number of prey items
131 (Krebs, 1999), ranging from 0 (all prey categories are equally present) to 1 (one category dominates
132 the community), the “Simpson” index $1 - D$ to estimate the dietary heterogeneity (from 0 to 1 as
133 complete evenness among prey categories), and the Shannon index: $H = \sum[(n_i/N) \times \ln(n_i/N)]$
134 to calculate the diversity from 0 with only a single prey category to high values with many prey
135 categories (Magurran, 2004). All these indices were determined using PAST (PAleontological
136 STatistics software, version 4.12; Hammer et al., 2001).

137 We created a Month \times Prey interaction matrix by grouping feeding records by month and
138 summing the number of each prey category. This matrix represents the total prey composition for

139 each survey month. The matrix was then converted into bipartite format and visualized using the
140 “bipartite” package in the R v3.5.2 (RStudio Team, 2018) to compare how prey contributions varied
141 among months (Dormann et al., 2009). Distinct colors were assigned to each month to aid
142 interpretation.

143 Dietary data on numbers of prey was employed for the Principal Coordinates Analysis
144 (PCoA). All prey items were extracted as numeric columns. A Bray–Curtis distance matrix was
145 computed in the “vegan” package in the PcoA (Oksanen et al., 2016). To visualize group structure,
146 we calculated group centroids and constructed convex hulls for each month. The relationships
147 between prey categories and diet months ordination were assessed, and only the strongest vector
148 loadings (top four) were projected onto the biplot. To test for differences in diet composition among
149 months, we conducted a permutational multivariate analysis of variance (PERMANOVA) with
150 4,999 permutations. Pairwise differences between months were assessed using pairwise
151 PERMANOVA on all month combinations. A significance level of different comparisons was
152 applied with P-value < 0.05.

153 Pianka’s index of niche overlap (Pianka, 1973) was used to calculate the trophic similarity or
154 segregation among surveyed months of the two seasons, by using the package “EcoSimR” set up
155 with 5000 replications and the “ra3” algorithm (Gotelli and Ellison, 2013), in the Rstudio:

156

$$O_{ij} = \frac{\sum_{n=1}^0 P_{ij} \times P_{ik}}{\sqrt{\sum_{n=1}^0 P_{ij}^2 \times P_{ik}^2}}$$

157 in which P_{ij} and P_{ik} are the proportions of number of food item i for the species categories j and
158 k , respectively.

159

RESULTS

160

161 *Diet Composition*

162 A total of 852 food items representing 25 prey categories were identified from the stomachs
163 of 128 individuals of *Cyrtodactylus leegrimeri* on Hon Khoai Island (Table 1). The diet primarily
164 consisted of small-sized invertebrates, encompassing 15 orders of the class Insecta and other taxa
165 belonging to Arachnida, Crustacea, and Annelida. It is noteworthy that we found attached plant
166 parts (including leaves) together with invertebrate prey, and shed skin of *C. leegrimeri* in the
167 stomach contents (Table 1).

168 Isoptera was identified as the primary prey category of *C. leegrimeri*, accounting for the
169 highest frequency of occurrence ($F = 0.41$, 27%), the largest number of prey items ($N = 670$,
170 78.6%), and the highest importance index ($I_x = 48.28\%$) (Table 1). Some prey groups also played
171 a vital role in the species' diet, including Orthoptera ($I_x = 16.62\%$), Lepidoptera (7.02%), Blattodea
172 (11.55%), and Araneae (7.68%). Remaining prey categories only occurred with small frequencies,
173 limited numbers, and low importance values (Table 1).

174

175 *Seasonal Variation*

176 The stomach contents of *C. leegrimeri* were collected in three different months and we
177 identified 298 prey items of 57 individuals in November 2015 of the rainy season, and 205 items
178 of 44 individuals in January 2016 and 349 items of 27 individuals in March 2025 of the dry season,
179 corresponding to the number of prey categories with 19, 18 and 11, respectively (Table 1; Fig. 2).
180 Isoptera consistently presented as the dominant prey group of *Cyrtodactylus leegrimeri* on the
181 island in all three months, exhibiting the highest frequency and abundance values (Fig. 3). The
182 importance index (I_x) score of Isoptera exceeded 35% in all periods, and even reached over 80%

183 in March 2025. During the rainy season (November 2015), Orthoptera, Lepidoptera, and Blattodea
184 were the following important prey groups (Ix values > 10%) (Table 1; Figs 2-3). During the dry
185 season, Orthoptera (Ix = 21.91%) and Blattodea (Ix = 17.28%) also represented key components
186 of the diet in January 2016, and Orthoptera (Ix = 18.27%) and Araneae (Ix = 12.93%) were
187 identified as major prey categories in March 2025 (Table 1; Figs 2-3).

188 In November 2015 (rainy season) and January 2016 (dry season), the estimated values of
189 Dominance_D and Simpson_1-D were approximately 0.5, indicating a moderate level of dietary
190 diversity and prey dominance. Similarly, the Shannon diversity index (Shannon_H) reflected a
191 medium level of prey richness, ranging from 1.28 to 1.33. However, these values were significantly
192 different from those recorded in March 2025 (dry season). In particular, the value of Dominance_D,
193 increased to a higher level, averaging 0.8 (95% CI: 0.74–0.84), while Simpson_1-D decreased to
194 less than half, averaging 0.199 (95% CI: 0.15–0.26) in March 2025. The Shannon_H index likewise
195 presented a threefold decline, averaging 0.55 (95% CI: 0.45–0.70), suggesting a reduction in the
196 prey richness, during this month (Table 2). The species exhibited a consistently narrow trophic
197 niche across the surveyed months, with low estimated niche breadth (B = 1.6–1.95) and
198 standardized values (< 0.06), indicating a highly selective feeding strategy (Table 2). Notably, the
199 standardized niche breadth in March 2025 was extremely lower than in the two remaining months
200 (Table 2).

201 Pianka's index revealed a relatively high degree of dietary overlap (P-value > 0.65) among
202 the survey months of the two seasons on Hon Khoai Island, indicating a similarity in the prey use
203 by *C. leegrimeri* (Fig. 4). The PCoA analysis showed that the first two axes accounted for 19.2%
204 (PCo1) and 14.9% (PCo2) of the variance in dietary composition of *C. leegrimeri* among the three
205 months. The PCoA ordination indicated a considerable overlap in the dietary variance space among

206 months and between the two seasons, with no significant differences (PERMANOVA, P-values >
207 0.05; Fig. 5). Based on the ordination of dietary space, Blattodea primarily contributed to the central
208 position observed in January 2016, Orthoptera predominantly to March 2025, and Isoptera to both
209 November 2015 and March 2025 (Fig. 5).

210

211 *Habitat Comparison*

212 We documented somewhat different patterns in the diet composition of *C. leegrimeri*
213 between forest and disturbed habitats. Specifically, two surveys in the sites of undisturbed forest,
214 recorded 19 (November 2015) and 18 prey categories (January 2016) (Fig. 1C), whereas the survey
215 conducted along road transects (March 2025; Fig. 1D), yielded only 11 prey categories. In disturbed
216 habitats, several prey groups, such as Diptera, Oniscomorpha, Cestoda, Opiliones, Gastropoda and
217 Insecta others, were not found in the stomach contents of *C. leegrimeri*. The Shannon diversity
218 index (Shannon_H) and the standardized dietary niche breadth declined by more than twice in the
219 disturbed habitats, but the dominance index (Dominance_D) increased (from 0.5 to 0.8) (Table 2).

220

221 DISCUSSION

222 Stomach contents were obtained from 128 individuals of *C. leegrimeri* during both the rainy
223 season (November 2015) and dry season (January 2016 and March 2025), identified with a total of
224 25 prey categories. The species presented a high degree of dietary specialization, with a high
225 dependence on Isoptera. The standardized niche breadth values were extremely narrow (< 0.06)
226 and the Shannon_H diversity values were also small (0.55–1.33), indicating that the species has a
227 narrow dietary range and exhibits selective feeding on specific prey categories. The results confirm
228 the central role of Isoptera in the trophic composition of *C. leegrimeri* on Hon Khoai Island. In

229 addition, the study identified other prey groups such as Orthoptera, Blattodea, Lepidoptera, and
230 Araneae as important dietary components for *Cyrtodactylus leegrimeri* on the Island. In natural
231 habitats, geckos have often been reported feeding on termites, spiders, grasshoppers, and ants (Vitt
232 and Zani, 1997; Vitt et al., 1997; Zamprogno and Teixeira, 1998; Colli et al., 2003; Bonfiglio et al.,
233 2006; Rocha and Anjos, 2007; Iturriaga and Marrero, 2013; Ngo et al., 2016a, 2024, 2025; Amadi
234 et al., 2020; Le et al., 2025).

235 Regarding the seasonal variation in the dietary composition, Isoptera was the most dominant
236 prey, and Pianka's index values indicating high dietary overlap among survey months and between
237 the two seasons. This pattern suggests that termites serve as a key food resource for *C. leegrimeri*,
238 due to their availability on Hon Khoai Island throughout both the rainy and dry seasons. The
239 reliance on ground-dwelling Isoptera may relate to the species' ground-foraging habit, as *C.*
240 *leegrimeri* was normally observed on the forest floor of soil substrate type or on trees at a low
241 height above the ground (Ngo, pers. obs.; Fig. 1). Another syntopic species, the Psychedelic rock
242 gecko *Cnemaspis psychedelica*, mainly occurring on rocky outcrops of the island (Ngo et al., 2018),
243 exhibited a higher consumption of Hymenoptera and Araneae (Ngo et al., 2024). For other tree or
244 ground-foraging lizard species, such as *Eutropis multifasciatus* (Ngo et al., 2014) and *Physignathus*
245 *cocincinus* (Nguyen et al., 2018), their food contents were dominated by Isoptera and Formicidae.
246 Meanwhile, rock-dwelling species, such as *Goniurosaurus catbaensis* and *G. huuliensis* were
247 found to feed predominantly on Arachnida and Orthoptera (Gryllidae), respectively (Ngo et al.,
248 2016a; Le et al., 2025). Ngo et al. (2025) also identified Araneae as the most important prey in the
249 diet composition of the rock-dwelling *Cyrtodactylus nigriocularis* in the southern mainland of
250 Vietnam. The differences suggest that the dietary partitioning between the ground+tree species
251 group (including, *C. leegrimeri*) and the rocky species group (*C. psychedelica*) might be

252 associated with preferred microhabitat use and foraging behavior. In particular, *C. psychedelica*, a
253 fast-moving and rock-dwelling species, prefers winged prey of Hymenoptera and Araneae, whereas
254 *C. leegrimeri*, a slower-moving and ground+tree-dwelling species, tends to exploit the abundant
255 termite colonies on the forest floor (Ngo et al., 2018, 2024; Ngo per. obs). Such niche partitioning
256 likely reduces interspecific competition for food resources, facilitating the coexistence and
257 abundance of both syntopic gecko species on the restricted area of Hon Khoai Island (Capitán et al.,
258 2015; Ngo et al., 2018, 2022).

259 The results revealed that human activities degrading the forest on Hon Khoai Island (Ngo et
260 al. 2018, 2022), are associated with the biased diet composition of *C. leegrimeri*. Although
261 termites (Isoptera) remained the dominant prey across all habitats, *C. leegrimeri* increasingly
262 relied on this prey group in disturbed habitats where overall prey richness was reduced. Unlike
263 groups absent from the diet contents of *C. leegrimeri* in the disturbed-open roads on Hon Khoai
264 Island (such as, Diptera, Oniscomorpha, Cestoda, Opiliones, and Gastropoda), termites characterize
265 a physiological thermal tolerance with complex nest-building behavior that buffers environmental
266 extremes, allowing them to persist across both thermally stable forest habitats and more variable
267 disturbed-open environments (Korb, 2003; Jones and Oldroyd, 2006; Woon et al., 2022). In
268 contrast, the groups rely more on intrinsic physiological flexibility (e.g., Diptera) or moisture-
269 dependent microhabitats (Cestoda, Oniscomorpha, Opiliones, Gastropoda), which generally
270 restricts them to more buffered forest conditions or limits their activity in disturbed, open habitats
271 with greater thermal and hydric stress (Edney, 1977; Willmer, 1982; Huey and Kingsolver, 2008).

272 In conclusion, there are no significant differences in the overall diet composition of *C.*
273 *leegrimeri* between seasons (rainy vs. dry) and between habitats (forest vs. road). However, our
274 findings recorded an increased reliance on termites and less prey richness in the disturbed habitats.

275 The shifts may result in the nutritional imbalance, and even pose a threat to the species if termite
276 availability declines. Together with the study of Ngo et al. (2024) on *Cnemaspis psychedelica*
277 (Endangered, IUCN Red List), we recommend protecting forest habitats. The protection is
278 therefore essential for the long-term persistence of *C. leegrimeri*, *C. psychedelica* and other
279 herpetofauna on Hon Khoai Island.

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416 **Tables.**

417 **Table 1.** Dietary composition of *Cyrtodactylus leegrimeri* between rainy and dry seasons, in forest and road habitats, on Hon Khoai

418 Island, Ca Mau Province (F – Frequency ; N – Number ; V – Volume ; Ix – Index of relative importance)

| | Rainy season | | | | Dry season | | | | | | | | Total | | | |
|------------------------|--------------------------|-----|---------|-------|-------------------------|-----|---------|-------|---------------------|-----|---------|-------|-------|-----|---------|-------|
| | November – 2015 (Forest) | | | | January – 2016 (Forest) | | | | March – 2025 (Road) | | | | | | | |
| | F | N | V | Ix | F | N | V | Ix | F | N | V | Ix | F | N | V | Ix |
| Araneae | 0.11 | 6 | 2292.37 | 8.16 | 0.07 | 3 | 273.42 | 3.43 | 0.33 | 9 | 226.39 | 12.93 | 0.14 | 18 | 2792.18 | 7.68 |
| Amblypygi | 0.02 | 1 | 183.34 | 1.02 | | | | | | | | | 0.01 | 1 | 183.34 | 0.45 |
| Opillions | 0.09 | 5 | 306.70 | 4.02 | 0.02 | 1 | 10.09 | 0.95 | | | | | 0.05 | 6 | 316.79 | 2.06 |
| Blattodea | 0.14 | 10 | 2433.47 | 10.02 | 0.32 | 16 | 1691.42 | 17.38 | 0.15 | 4 | 268.32 | 6.45 | 0.20 | 30 | 4393.21 | 11.55 |
| Coleoptera | 0.02 | 1 | 8.94 | 0.71 | 0.02 | 1 | 100.93 | 1.17 | 0.07 | 2 | 6.57 | 2.69 | 0.03 | 4 | 116.44 | 1.29 |
| Diptera | 0.02 | 1 | 4.35 | 0.70 | 0.02 | 1 | 2.28 | 0.93 | | | | | 0.02 | 2 | 6.63 | 0.60 |
| Ephemeroptera | | | | | | | | | 0.04 | 1 | 13.65 | 1.39 | 0.01 | 1 | 13.65 | 0.31 |
| Hemiptera | 0.04 | 2 | 58.91 | 1.50 | 0.02 | 1 | 25.61 | 0.98 | 0.04 | 1 | 257.57 | 2.42 | 0.03 | 4 | 342.09 | 1.48 |
| Hymenoptera | 0.02 | 2 | 20.58 | 0.84 | | | | | | | | | 0.01 | 2 | 20.58 | 0.36 |
| Hymenoptera-Formicidae | 0.09 | 26 | 78.48 | 5.97 | 0.02 | 1 | 0.64 | 0.92 | 0.04 | 1 | 0.24 | 1.33 | 0.05 | 28 | 79.35 | 2.98 |
| Isoptera | 0.32 | 211 | 1974.41 | 37.56 | 0.30 | 147 | 2233.43 | 39.26 | 0.81 | 312 | 5861.95 | 81.73 | 0.41 | 670 | 10069.8 | 48.28 |
| Lepidoptera | 0.02 | 2 | 5331.96 | 10.07 | 0.02 | 1 | 684.49 | 2.61 | 0.15 | 4 | 292.79 | 6.56 | 0.05 | 7 | 6309.23 | 7.02 |
| Orthoptera | 0.09 | 5 | 4692.43 | 11.63 | 0.11 | 6 | 6953.94 | 21.91 | 0.41 | 12 | 839.44 | 18.27 | 0.16 | 23 | 12485.8 | 16.62 |
| Thysanoptera | | | | | 0.05 | 2 | 262.18 | 2.49 | | | | | 0.02 | 2 | 262.18 | 0.81 |
| Thysanura | | | | | 0.07 | 13 | 32.88 | 4.47 | | | | | 0.02 | 13 | 32.88 | 1.32 |
| Insect - others | 0.05 | 3 | 219.21 | 2.47 | 0.07 | 3 | 16.87 | 2.80 | | | | | 0.05 | 6 | 236.08 | 1.99 |
| Insect larva | 0.02 | 1 | 26.70 | 0.74 | | | | | | | | | 0.01 | 1 | 26.70 | 0.32 |
| Isopoda | | | | | 0.05 | 4 | 43.24 | 2.27 | 0.07 | 2 | 110.91 | 3.13 | 0.03 | 6 | 154.14 | 1.40 |

| | | | | | | | | | | | | | | | | |
|-------------------|-----------|------------|--------------|---------------|--------------|------------|--------------|---------------|--------------|------------|----------------|------|---------------|------------|---------|------|
| Oniscomorpha | 0.11 | 6 | 46.47 | 4.26 | 0.02 | 1 | 26.13 | 0.98 | | | | | 0.05 | 7 | 72.61 | 2.16 |
| Polydesmoidea | 0.02 | 1 | 82.38 | 0.84 | | | | | | | | | 0.01 | 1 | 82.38 | 0.37 |
| Lumbricina | | | | | 0.02 | 1 | 601.52 | 2.40 | | | | | 0.01 | 1 | 601.52 | 0.79 |
| Polydesmida | | | | | | | | | 0.04 | 1 | 11.48 | 1.38 | 0.01 | 1 | 11.48 | 0.31 |
| Scolopendromorpha | 0.02 | 1 | 42.90 | 0.77 | | | | | | | | | 0.01 | 1 | 42.90 | 0.33 |
| Cestoda | 0.16 | 9 | 1159.92 | 8.28 | 0.05 | 2 | 526.61 | 3.14 | | | | | 0.09 | 11 | 1686.53 | 4.68 |
| Gastropoda | 0.07 | 5 | 229.02 | 3.30 | 0.02 | 1 | 32.38 | 1.00 | | | | | 0.04 | 6 | 261.40 | 1.75 |
| Plant parts | 0.01 | 1 | | | | | | | | | | | 0.01 | 1 | | |
| Shed skin | | | | | | | | | 0.01 | 1 | | | 0.01 | 1 | | |
| Total | 57 | 298 | 19193 | 112.87 | 44.00 | 205 | 13518 | 109.09 | 27.00 | 349 | 7889.30 | | 128.00 | 852 | 40599.9 | |

419

420 **Table 2.** Dominance (D), Simpson (1-D), Shannon indices (H) and Niche breadth of *Cyrtodactylus*
 421 *leegrimeri*, among three survey months, in forest and road habitats on Hon Khoai Island.

| | November (Forest) | | | January (Forest) | | | March (Road) | | |
|-------------------------------|-------------------|--------------|--------------|------------------|--------------|--------------|--------------|--------------|--------------|
| | | 95% Lower | 95% Upper | | 95% Lower | 95% Upper | | 95% Lower | 95% Upper |
| Taxa_S | 19 | | | 18 | | | 11 | | |
| Individuals | 298 | | | 205 | | | 349 | | |
| Dominance_ D | 0.51 | 0.44 | 0.57 | 0.52 | 0.43 | 0.59 | 0.80 | 0.74 | 0.84 |
| Simpson_1-D | 0.49 | 0.43 | 0.56 | 0.48 | 0.41 | 0.57 | 0.19 | 0.16 | 0.26 |
| Shannon_H | 1.33 | 1.18 | 1.52 | 1.28 | 1.11 | 1.51 | 0.55 | 0.45 | 0.70 |
| Niche breadth | 1.95 | | | 1.89 | | | 1.61 | | |
| Standardized Niche breadth | 0.05 | | | 0.05 | | | 0.02 | | |

422

423 **Legends of figures**

424 **Fig. 1.** *Cyrtodactylus leegrimeri* on a tree (A), on the ground (B), and habitat types of (C) forest
425 (D) road on Hon Khoai Island

426 **Fig. 2.** Frequency; Number; and Importance index of prey categories consumed by *Cyrtodactylus*
427 *leegrimeri* during survey months, on Hon Khoai Island

428 **Fig. 3.** The Bipartite network for prey contributions of *Cyrtodactylus leegrimeri* varied among
429 months

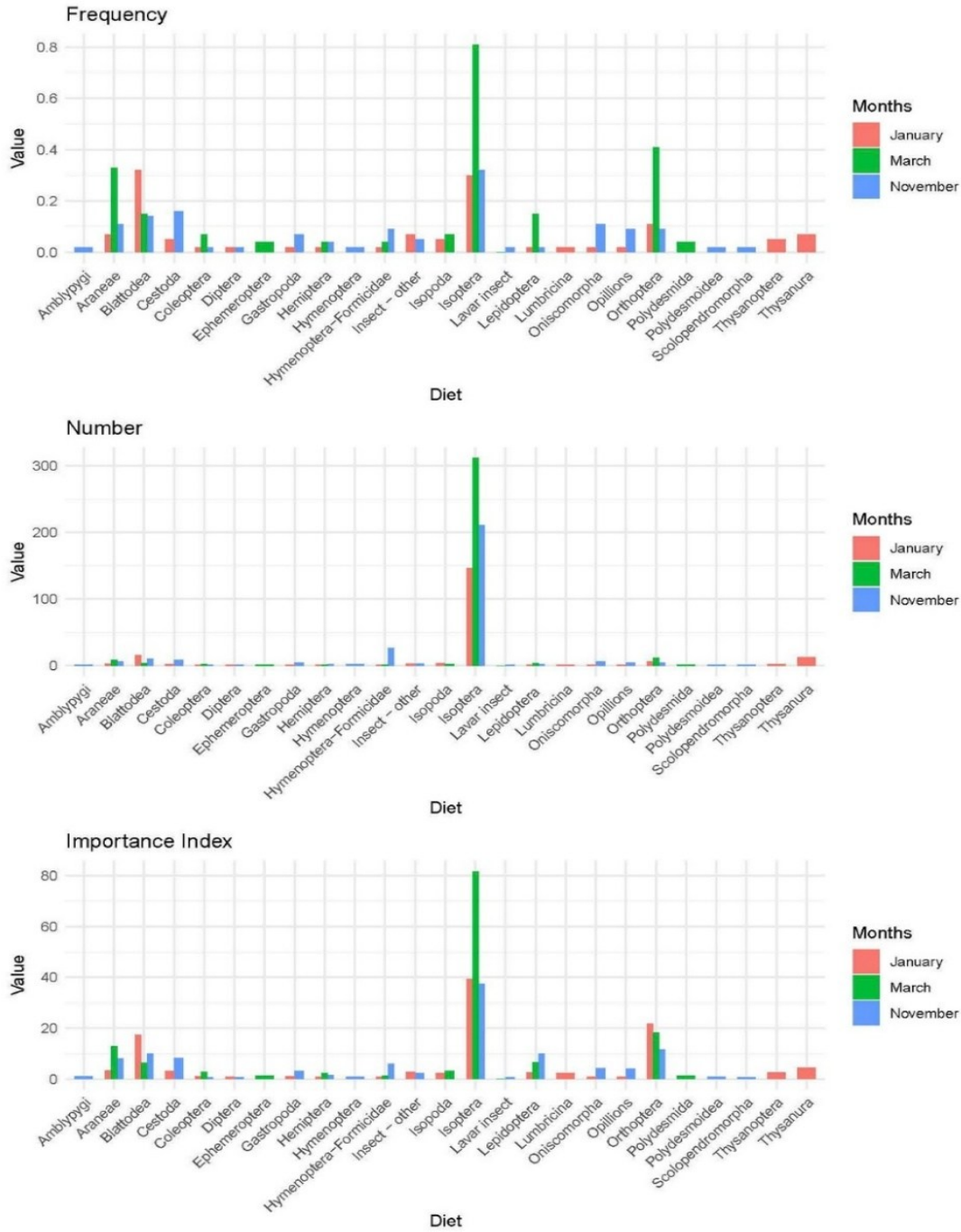
430 **Fig. 4.** The simulated Pianka's index of trophic niche overlap among survey months of
431 *Cyrtodactylus leegrimeri* on Hon Khoai Island (The red line indicates the observed overlap).

432 **Fig. 5.** The PcoA comparison of diet niches of *Cyrtodactylus leegrimeri* among three month
433 surveys, with contribution of strongest prey categories, explained by axes of principal components
434 (PCo1 and PCo2).



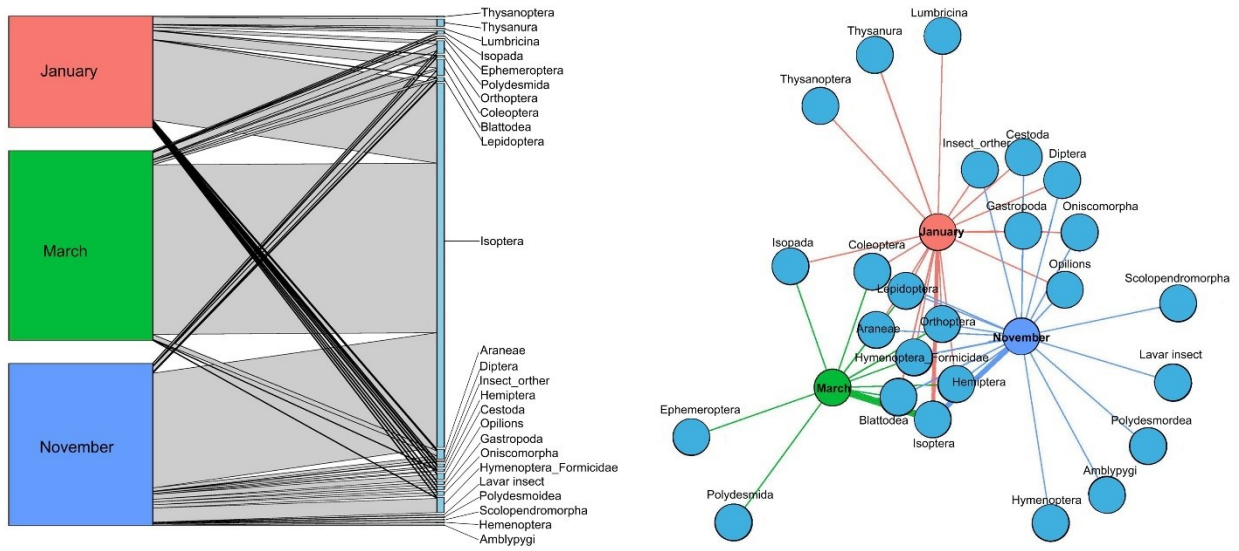
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436 **Fig. 1.**



437

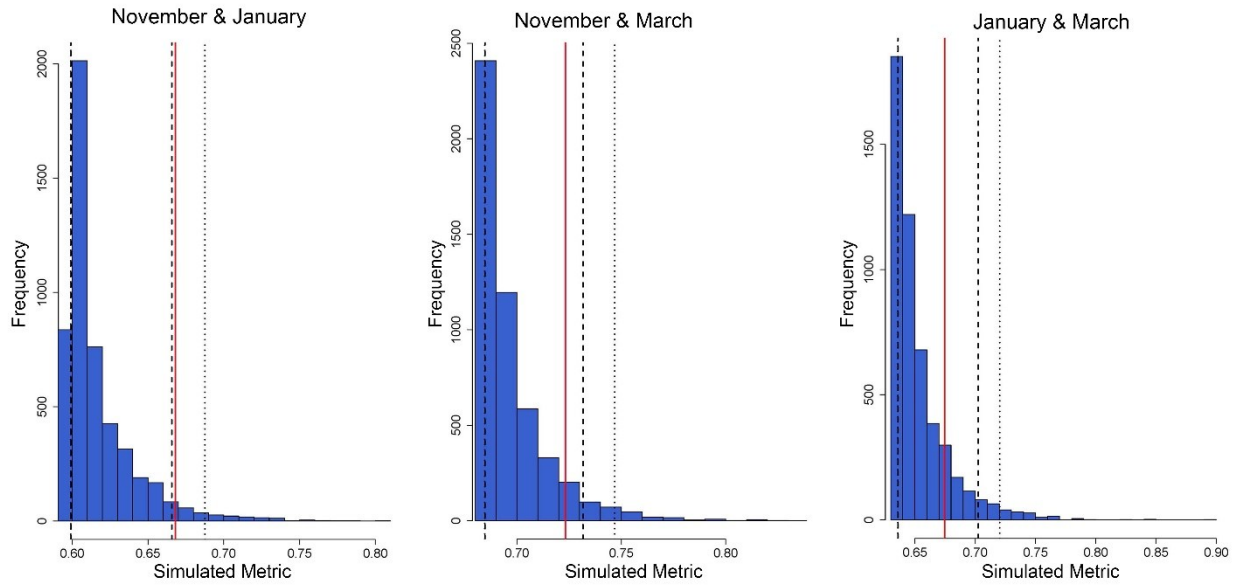
438 Fig. 2.



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440 **Fig. 3.**

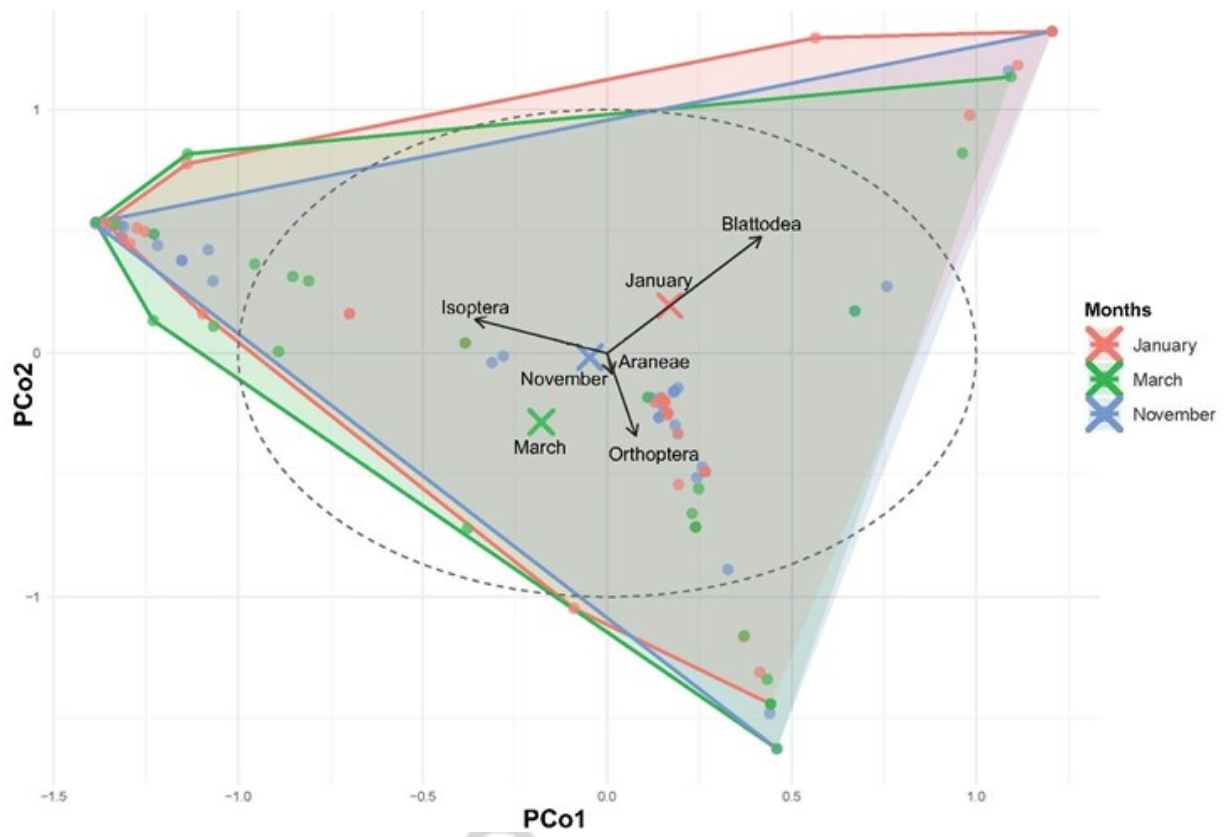
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442 **Fig. 4.**

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444 Fig. 5.

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