

Diversity and efficacy of insect sampling methods in an urban tropical dry forest of the Colombian Caribbean (#99645)

1

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Diversity and efficacy of insect sampling methods in an urban tropical dry forest of the Colombian Caribbean

Rodrigo Sarmiento-Garcés¹, Daniel Posada-Echeverría¹, Rafik Neme^{Corresp. 1}

¹ Departamento de Química y Biología, Universidad del Norte, Barranquilla, Atlántico, Colombia

Corresponding Author: Rafik Neme
Email address: rneme@uninorte.edu.co

Tropical Dry Forests (TDFs) are among the most threatened ecosystems in Colombia and the world. Characterized by distinct dry and wet seasons, TDFs support a wide array of flora and fauna and offer crucial ecosystem services, despite facing ongoing threats like deforestation and climate change. This study provides a comprehensive comparison of three insect sampling methods within an urban TDF area in Puerto Colombia, Atlántico, in the Colombian Caribbean. We assessed the efficacy of Malaise traps, pitfall traps, and canopy fogging in capturing insect diversity and examined the presence of various trophic guilds. In total, we collected 3135 insects, comprising 413 morphospecies. The results indicate Malaise traps as the most effective, capturing a diverse range of species, followed by pitfall traps, while canopy fogging yield the lowest richness. Each method exhibits unique advantages and captures specific species, revealing that the methods complement each other in understanding insect diversity in these fragile and understudied ecosystems.

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2 **Colombian Caribbean**

3
4 **Abstract**

5
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15 yield the lowest richness. Each method exhibits unique advantages and captures specific
16 species, revealing that the methods complement each other in understanding insect diversity in
17 these fragile and understudied ecosystems.
18

19

20 **Introduction**

21

22 Tropical dry forests (TDF) occur in dryland environments in regions with tropical climate. In the
23 neotropics they are defined as a biome which encompasses a set of floristic groups, with a wide
24 distribution ranging from Mexico to Argentina (DRYFLOR et al., 2016). TDFs hold significant
25 value due to their strong seasonality in precipitation, and marked heterogeneity in rainfall
26 seasonality, soil characteristics, frost, and altitude (González-M et al., 2018). TDFs are vital for
27 biodiversity conservation and provide critical ecosystem services, including carbon storage,
28 climate regulation, nutrient cycling, economic and recreational values (Lewis et al., 2009; Siyum,
29 2020; Vinya et al., 2019; Zhou et al., 2013). However, they are among the most threatened
30 ecosystems globally, suffering from deforestation, land-use change, and climate change,
31 making them the one of the most threatened and impacted tropical terrestrial ecosystem
32 (Ferrer-Paris et al., 2019; Miles et al., 2006).
33

34

35 In the Colombian Caribbean, TDFs have been greatly impacted by human activities, as a
36 consequence of fragments being located on lands with the most fertile soils which leads to the
37 promotion of productive activities, leading to an increase of land-use change for agricultural, and
38 urbanization (Correa Ayram et al., 2020). The remaining dry forest patches are highly
39 fragmented, leading to the isolation of plant and animal populations and reducing their resilience
40 to environmental stressors (Pizano & Garcia 2014).

41

42 Despite the singularity, importance, and degree of threat to Neotropical dry forests, studies on
43 them are scarce and limited to a few biotic communities (IAvH. 1998, Pizano & Garcia 2014). In
44 the case of insects, research has primarily focused on specific taxa, with limited studies
45 covering the entire terrestrial insect community or a broad spectrum of trophic levels (Aldana-
46 Domínguez et al 2017).

47

48 Insects play critical roles in TDFs, participating in processes like pollination, nutrient cycling,
49 decomposition, and providing ecosystem services, including pollination and pest control.
50 Understanding the diversity, community structure, and trophic guilds of insects in these forests
is crucial for effective conservation measures. **There is a need for more comprehensive**

51 research on the use of Malaise traps, a tool for capturing a wide variety of insect species, in
52 specific regions like Colombia (Hausmann et al., 2020).

53

54 Various insect sampling methods, each with its limitations and biases, are essential tools for
55 studying insect diversity and community structure amidst the ongoing challenges of mass
56 extinctions and climate change. While each method has strengths and limitations, using multiple
57 approaches, including Malaise traps, pitfall traps, and canopy fumigation (fogging), can offer a
58 more comprehensive understanding of insect diversity and community structure in ecosystems.
59 Malaise traps have proven effective for capturing flying insects and have been crucial for
60 capturing a wide range of taxa and trophic guilds, providing valuable insights into the diversity
61 and community structure of insects in TDFs (Borkent et al., 2018; Hausmann et al., 2020). In
62 this study, we assessed the diversity and trophic guilds present in a tropical dry forest of Puerto
63 Colombia, evaluating the effectiveness of these three commonly used sampling methods.

64

65 **Methods**

66

67 **Area of study**

68

69 Our study was conducted in the municipality of Puerto Colombia, Atlántico, part of the
70 Metropolitan Area of Barranquilla, in the Colombian Caribbean Region. The area of interest is in
71 the Pericaribbean Arid Belt (Hernández et al., 1992), with patches of Tropical Dry Forest (TDF)
72 (Holdridge, 1971, Sánchez-Azofeifa et al., 2005). It has an altitude of 4 meters above sea level,
73 an approximate rainfall of 900mm, an average temperature of 27°C, relative humidity ranging
74 between 75 - 85%, and evaporation rates of 1500 to 1900 mm (IDEAM 2020).

75

76 The sampling was carried out in a relic of Tropical Dry Forest inside Universidad del Norte in
77 November 2019 with an approximate area of 8.5 Ha, surrounded by a mosaic of developing
78 urban landscapes dominated by secondary vegetation. The forest presents a well-defined
79 stratification with a low or herbaceous stratum dominated mainly by herbs and creeping lianas,
80 a medium or shrubby stratum composed of shrubs or small trees (DBH \leq 5 cm), and a canopy
81 or tree stratum made up of larger trees (DBH \geq 5 cm) reaching heights of up to approximately
82 12 meters.

83

84 **Sampling of terrestrial entomofauna**

85

86 Different collection methods were used to obtain a representative sample of the available
87 entomofauna in the forest described above. The effectiveness of three collection methods was
88 evaluated, and the contribution and complementarity of each method to the global collection
89 were assessed. The methods used were selected to sample each of the strata present in the
90 forest. For this purpose, pitfall traps were used for the low stratum, Malaise-type flight
91 interception traps for the medium or shrubby stratum, and the fumigation method of the canopy
92 or fogging for collections in the high or arboreal stratum. The collections were carried out in
93 October 2019 during the season of humidity and maximum temperature values. Below is a brief
94 description of the collection methods used.

95

96 **Pitfall traps**

97

98 To capture terrestrial insects, present on the soil surface, a transect of 12 pitfall traps was
99 installed, 20 meters between traps. The period of activity per trap was eight days, for a total
100 exposure of 192 hours per trap. Each trap consisted of a plastic container (25 cm in diameter,

101 25 cm deep), placed at ground level without any type of attractant, half-filled with a mixture of
102 water, detergent, and 96% alcohol.

103

104 Malaise traps

105

106 In the case of insects associated with the herbaceous and shrubby or medium stratum, four
107 Malaise-type flight interceptor traps were installed, separated by 30 to 50 meters between each
108 trap. The installed traps consisted of the classic Malaise traps with front height dimensions of
109 1.80 meters, rear height of 1.10 meters, and a length of 1.60 meters. The collection devices
110 were active for eight days (192 hours per trap).

111

112 Canopy fogging

113

114 For the canopy stratum, an area of 30 m² was fumigated with broad-spectrum pyrethroid
115 insecticide (Pyrethrins 3% w/v + Piperonyl Butoxide 15% w/v EW), from which four independent
116 samples of 4.2 m² were taken (1.2m x 3.5m). The collection exposure time after fumigation was
117 two hours, during which the insecticide had a residual effect.

118

119 All the collection material was labeled and preserved in 96% alcohol until further processing.

120

121 Taxonomic identification

122

123 Once the material was collected from each of the methods, taxonomic determination and
124 quantification were carried out. For each collection method, all the organisms were separated at
125 the morphospecies level and identified to the lowest possible taxonomic level that would allow
126 the identification of the trophic guild. The identified morphospecies were counted to establish
127 their abundances. The keys and diagnosis of González & Carrejo (1992), White (1998), Arnett y
128 Thomas (2000), Triplehorn & Johnson (2005), Fernández y Sharkey (2006) were used for the
129 taxonomic determination at the family and superfamily level.

130

131 Allocation of trophic guilds

132

133 For the characterization of the trophic groups of each taxon, a review of the literature on their
134 life histories and eating habits was carried out based on the literature indicated for taxonomic
135 identification. Thus, 11 categories or trophic guilds were established: omnivorous, parasitoids,
136 mycophages, saprophagous/coprophagous, nectarivorous/polynivorous, gall-forming
137 phytophagous, chewing phytophagous, sap-sucking phytophagous, xylophagous/stem-
138 boring/seed-eating phytophagous, predators, and hematophagous. This classification was
139 based on the definition of functional groups by Steneck (2000) and Clavijo-Awazacko &
140 Amarillo-Suárez (2013), who define them as a group of species that play equivalent functional
141 roles in communities and ecosystems.

142

143

144 Data analysis

145

146 The insect communities present in each collection method were compared using confidence
147 intervals (95%) (Chao1 estimator) of the rarefaction curves and extrapolation of the Hill
148 numbers, using the richness (q=0) and diversity values (q=1 and q=2) (Chao et al., 2014, Hsieh
149 et al., 2016) with a bootstrap of 999 iterations. Likewise, through these estimators, the
150 percentage of sampling coverage was evaluated for each of the methods used, as well as for
151 each of the orders of insects collected, evaluating the collection methods.

152

153 For the comparison of the composition between communities according to each collection
154 method (Beta diversity), we used a multivariate analysis of variance (Permanova), using the
155 adonis function, with 999 permutations in the vegan package (Oksanen et al., 2017). With the
156 betapart package (Baselga & Orme, 2012), diversity was partitioned into nestedness and
157 turnover to understand if the result of dissimilarity is due to substitution or loss of species
158 between the different collection methods.

159

160 For the analysis of trophic guilds, a one-way ANOVA was performed to compare the richness of
161 trophic guilds for each collection method. Additionally, the Equity index J was estimated to
162 evaluate the distribution of richness and understand if there is dominance by a few guilds in
163 some of the methods.

164

165 All statistical analyses were performed using the statistical environment R (R Core Team, 2023).

166

167 Results

168

169 A total of 3135 individuals of the class Insecta, distributed among 413 morphospecies, were
170 collected. The orders with the highest representation were Hymenoptera (137 sp, 1280
171 individuals), Coleoptera (125 sp, 1237 individuals), Diptera (87 sp, 377 individuals), and
172 Hemiptera (36 sp, 120 individuals), which represent about 95% of the total reported richness.
173 The other orders had considerably lower presence with fewer than nine species (Table
174 1)(Figure 1).

175

176 Of all the species captured, the Malaise trap was the most effective capture method (284 sp,
177 1174 individuals), presenting more than twice the species richness than the second most
178 effective method (104 sp, 1127 individuals), the pitfall traps. Finally, canopy fogging was the
179 method that presented the lowest richness values (78 sp, 834 individuals), presenting about a
180 third of the richness collected by the Malaise method.

181

182 The Malaise trap was mainly represented by the orders Hymenoptera, Diptera, and Coleoptera,
183 which accounted for 91% of the species richness collected for this method. The pitfall traps
184 mainly collected organisms from the orders Coleoptera and Hymenoptera (mainly ants) with
185 72% of the species present, while the canopy fogging method was represented mostly by the
186 orders Hymenoptera, Coleoptera, and Hemiptera with 82% of the species (Figure 1)(Figure 2).

187

188 The sampling coverage was adequate for each of the collection methods, where the canopy
189 fogging and pitfall trap methods had the best performance, collecting about 95% of the potential
190 species compared to the Malaise trap method, which had a coverage percentage of 85%
191 (Figure 2) (Figure 3). However, despite the good results found in sampling coverage, when
192 insect orders are analyzed separately, we see a somewhat different picture, where orders such
193 as Mantodea and Orthoptera present percentages of sampling coverage below 60%. The
194 remaining orders had a representation greater than 80% coverage percentage (Supplemental
195 Figure 1). Orders such as Collembola, Embioptera, Neuroptera, and Zygentoma were not
196 included in this analysis due to their low representativeness with abundances less than five
197 individuals, which denotes the low effectiveness of these methods for the collection of the
198 orders.

199

200 The rarefaction curves showed significant differences in richness for each of the methods
201 explored, with non-overlapping confidence intervals. The Malaise trap presented close to 70%
202 of the total collected richness, followed by the pitfall traps with 25% of the richness, and the

203 canopy spraying method with 20%. This result shows us that Malaise traps are the most
204 effective mechanism for collecting the terrestrial insect community, with the ability to capture
205 almost three times more richness than the other two methods compared (Figure 3A) (Figure
206 4A).

207
208 Diversity analysis, assuming relative abundance, showed similar results to those obtained in
209 richness analyses, where the Malaise trap presents the highest diversity values for both typical
210 species (Shannon, $q=1$) and abundant species (Simpson, $q=2$) (Figure 3B and 3C) (Figure 4B
211 and 4C).

212
213 For each type of trap, we can see that the typical species (Shannon, $q=1$) in the Malaise trap
214 occupy about 42% of the richness, while the abundant species (Simpson, $q=2$) are 20% of all
215 the species. The pitfall traps had a percentage of 20% of typical species, while the abundant
216 species were 11%. The canopy fogging method reported 24% typical species and 15%
217 abundant species. The previous results show us that, despite the high efficiency of the traps,
218 the collected communities are dominated by a low number of species, which tells us of a low
219 equality in the distribution of abundance.

220
221
222 Among the unique features of each trap (Figure 4A-C) (Figure 5A-C), we can see that the
223 canopy fogging method provides 51 species of the families Buprestidae, Cassidinae,
224 Cerambycidae, Chrysomelidae, Cicadellidae, Coccinellidae, Corylophidae, Curcuionidae,
225 Dermestidae, Dictyopharidae, Formicidae, Histeridae, Homoptera, Lampyridae, Mantidae,
226 Membracidae, Miridae, Ploiarridae, Reduvidae, Thespidae, Tingidae, and a high richness of
227 Psocoptera.

228
229 For its part, pitfall traps presented 74 exclusive species of the families Acanthopodiae,
230 Aphodiinae, Aradidae, Blattellidae, Blattidae, Bolboceratinae, Ceratocanthidae, Chrysomellidae,
231 Cicadellidae, Curculionidae, Elateridae, Endomychidae, Formicidae, Gryllidae, Histeridae,
232 Isotomidae, Lygaeidae, Muscidae, Mutilidae, Mycetophilidae, Nicoletiidae, Phoridae,
233 Pompilidae, Pyrrhocoridae, Scarabeidae, Silvanidae, Staphylinidae, and Tenebrionidae.

234
235 Finally, the Malaise traps presented 241 exclusive species of the family Agromyzidae,
236 Ampulicidae, Anisopodidae, Anobiidae, Anthicidae, Anthomyiidae, Apoidea, Asilidae,
237 Bostrichidae, Braconidae, Bradynobaenidae, Calliphoridae, Canacidae, Cassidinae,
238 Cecidomyiidae, Cerambycidae, Ceratopogonidae, Cercopidae, Chalcidoidea, Chamamyiidae,
239 Chironomidae, Chloropidae, Chrysidoidea, Chrysomelidae, Cicadellidae, Cleridae, and
240 Clusiidae.

241
242 In terms of the composition of the species and their abundance (Bray-Curtis dissimilarity), we
243 found that there were significant differences between each of the collection methods ($F=20.10$;
244 $P < 0.001$). This shows us that, in addition to the differences in richness and diversity between
245 methods, each of the methods collects a particular composition of fauna. Now, it is important to
246 understand whether these dissimilarity results are due to subsets between samples
247 (nestedness) or due to a high degree of species turnover (turnover). The results show, first of
248 all, that there is a high dissimilarity between samples ($SOR=0.88$), which is given almost entirely
249 by a high degree of species turnover ($SIM=0.77$; 87%) and to a lesser degree by loss or subsets
250 of species (nestedness) ($SNE=0.11$; 13%) (Figure 4D-E)(Figure 5 D-E). This is reflected in the
251 fact that each trap has a high percentage of unique species, where the canopy fogging presents
252 65% of exclusive species, the pitfall 71%, and the Malaise 85%.

253

254 These results indicate that although methods such as pitfall traps or the canopy fogging method
255 have lower species richness, they enable access to species composition that cannot be
256 obtained by other methods.

257
258

259 When we observe the dissimilarity between paired samples, we can see that in general, all the
260 samples share few elements (Figure 5D, SOR = 0.86 - 0.89). However, the greatest distance in
261 terms of composition is between the pitfall traps and the canopy fogging method, where there is
262 a very low degree of nesting (nestedness) (SNE= 0.02). For its part, the flight interception traps,
263 compared to the canopy fogging method, had a greater number of nested or shared species,
264 which shows us that although with specific elements, the Malaise trap can collect some of the
265 elements collected by the fogging method (sne= 0.21).

266

267 In terms of the representation of trophic guilds, we were able to identify the presence of 11
268 functional groups (Figure 6-7). The Malaise traps were able to collect all guilds, while the pitfall
269 and canopy fogging traps collected nine of them. As with species richness, the difference in
270 trophic guilds for each collection method was also significant ($F=31$, $P<0.001$), with the Malaise
271 trap having a greater ability to collect trophic guilds (Figure 6-7).

272

273 When we analyzed the affinity of each collection method for each trophic guild, considering the
274 number of species, we observed that in general, the traps had a high evenness (J) with values
275 greater than 80% (Table 1). This means that each collection method did not show dominance by
276 a few trophic guilds. However, despite these results, we can see some trends worth highlighting.
277 The Malaise traps had a greater representation of the saprophytic/saprophagous/coprophagous
278 guilds (22%), parasitoids (21%), and predators (20%). Meanwhile, in the pitfall traps, their
279 greater representation was given by the saprophytic/saprophagous/coprophagous (26%),
280 omnivores (20%), and predators (16%). Finally, in the canopy fogging traps, the highest
281 representation was given by the omnivorous (17%), parasitoid (16%), predatory (15%), and
282 phytophagous sap-sucking guilds (14%) (Figure 7).

283

284

285

286 Discussion

287

288 We identified a rich diversity of insects in a relict of dry forest in Puerto Colombia, within the
289 Metropolitan area of Barranquilla, with sampling coverage values exceeding 85%. Our collection
290 methods not only complement each other but also exhibit a high turnover of species. The
291 Malaise traps were notably effective, securing more than double the species compared to the
292 second most effective method, pitfall traps.

293

294 Despite the forest's size and context, it sustains various trophic guilds, indicating a valuable
295 provision of ecosystem services offered by conserving such urban forest remnants. The
296 entomofauna in our study area demonstrated a richness either equivalent or superior to those
297 reported in other continental dry forests (Janzen 1968, 1973a, 1973b; Vasconcellos et al. 2010;
298 Macedo Reis et al. 2019; Silva et al. 2017) and Caribbean islands (Janzen 1968, 1973a,
299 1973b). This richness, seen even in a limited area, highlights the conservation of a significant
300 number of species and trophic guilds due to the presence of defined stratification. The diversity
301 in terrestrial ecosystems correlates with tree abundance and is influenced by both structural
302 complexity and plant species richness (McKinney, 2008; Helden et al, 2012).

303

304 When our results are compared to larger-scale studies (Janzen 1968), those incorporating
305 various time windows (Janzen 1968, Silva et al. 2017), or those at the molecular scale (Janzen
306 et al. 2009), the depicted diversity is but a fraction of what tropical dry forests can potentially
307 harbor. Our data collection was temporally limited (wet season), and the reliance on
308 morphological characters for morphospecies separation might have underestimated the
309 richness within some species groups, particularly those with cryptic species (Janzen et al. 2009;
310 Yu et al. 2012; Geiger et al. 2016).

311
312 It's crucial to acknowledge that our comparative global data derives from studies utilizing
313 different collection methods or focusing on specific trophic guilds (Janzen 1968, 1973a, 1973b;
314 Vasconcellos et al. 2010; Macedo Reis et al. 2019; Silva et al. 2017). There is a noticeable
315 scarcity of broad-spectrum collection methods aimed at diverse taxa and trophic guilds in the
316 Caribbean.

317
318 In our study, the Malaise trap demonstrated the highest efficiency in insect collection, supporting
319 findings documented across various ecosystems (Faulds & Crabtree 1995; Darling & Packer
320 1988; Shweta & Rajmohana 2016; Thomas & Sheikh 2016). The trap efficiently captures a wide
321 range of terrestrial insects, collecting both ground-moving organisms and those flying at low
322 levels (approximately 2.5 meters) (Marston 1965; Thomas & Sheikh 2016).

323
324 Secondary environments like ours possess undergrowth plant biomass providing abundant
325 resources for insects, facilitated by tree species adapted to water deficits with smaller canopies,
326 allowing sunlight penetration and undergrowth development (Herazo-Vitola et al. 2016). This
327 environment favors both undergrowth individuals and associated entomofauna during various
328 successional stages.

329
330 The Malaise trap is not only adept at collecting a plethora of insect orders with special affinity for
331 the most diverse ones—Coleoptera, Hymenoptera, and Diptera (Triplehorn & Johnson 2005)—
332 but also effectively doubles the diversity values reported by other methods.

333
334 Pitfall traps, while popular, inherently favor certain taxa, limiting their functional scope to ground-
335 dwelling species like Tenebrionids, Staphylinids, Scarabeinos, Formicidids, and Blattidids
336 (Montgomery et al. 2021). Contrastingly, canopy fogging, a general inventory method with a
337 wide taxonomic scale (Basset et al. 1997), yielded lower values than Malaise traps and did not
338 significantly differ from pitfall traps. This outcome, surprising given previous studies (Erwin
339 1983; Adis et al. 1984), mirrored the richness observed in the canopies of other continental dry
340 forests (Neves et al. 2010; Diodato & Fuster 2016), showcasing the variability in canopy habitats
341 under different pressures or stresses (Adis et al. 1984).

342
343 The Malaise trap excelled not only in species richness but also in abundance distribution across
344 species ($q=1$ and $q=2$), whereas pitfall and fogging methods exhibited narrower affinities for
345 select species. Such values may correlate with the habitat dynamics where collections occurred,
346 given the forest's successional state offering a stable microclimate and abundant resources
347 within the medium and creeping strata where Malaise traps operate.

348
349 Furthermore, methodological differences were not confined to richness and evenness; there
350 were distinct variations in species identity found, underscoring the specificity of each method.
351 This variance is likely due to the distribution of insect guilds across forest strata (McKinney,
352 2008; Helden et al., 2014), as our intervened forest can host well-differentiated taxa layers.

353

354 Guild diversity, akin to taxonomic diversity, was unique for each method regarding both diversity
355 measures and dominant guild identity per trap type. For instance, Malaise traps predominantly
356 captured saprophytes/saprophages/coprophages due to the presence of dipterans (Montgomery
357 et al., 2021), which, owing to their flight habits, rarely appear in pitfall traps. The traps also
358 registered predators from the Hymenoptera, Coleoptera, and Diptera families, capitalizing on
359 prey available on herbaceous and shrubby foliage (Triplehorn & Johnson 2005), and parasitoids
360 employing similar foraging techniques (Shweta & Rajmohana, 2016).

361
362 Pitfall traps primarily recorded saprophytic/saprophagous/coprophagous and omnivorous guilds,
363 with decomposers mostly belonging to the Coleoptera order exploiting soil detritus (Nichols et
364 al., 2008). Predatory guilds, represented by Coleoptera and Hymenoptera (Formicidae), utilized
365 herbaceous vegetation and creeping lianas as major resources (Janzen, 1973b).

366
367 Lastly, canopy fogging revealed four trophic guilds with comparable richness values, all
368 exploiting sprouts, flowers, or fruits in tree crowns, interacting with associated prey (McKinney,
369 2008; Helden et al., 2012). This complex interaction of species and guilds underlines the
370 intricate biodiversity sustained in these crucial urban forest remnants.

371

372 **Concluding remarks**

373

374 Our findings underscore the importance of employing multiple insect sampling methods to attain
375 a comprehensive understanding of the insect diversity and community structure within TDFs.
376 The unique attributes of each sampling technique are pivotal in capturing different insect taxa
377 and trophic guilds, highlighting their complementary roles. The rich biodiversity found within the
378 urban TDF of Puerto Colombia, even in the face of environmental stressors, emphasizes the
379 necessity for concerted conservation efforts. This research not only contributes to the limited
380 data on insect communities in the Caribbean TDFs but also supports global initiatives aimed at
381 biodiversity conservation in rapidly urbanizing environments.

382

383 **Acknowledgements**

384

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386 establishment of this project as a Partner Group. We thank Cecilia Torres, Andres de la Hoz,
387 and Anahi Barros for their help in the field. We thank Universidad del Norte for allowing us
388 access to the relict of forest. We hope projects like this will help the preservation of this patch of
389 land for further scientific use.

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Figure 1

Richness by orders of insects and efficiency of three different collection methods and their efficiency in a relict of dry forest in the municipality of Puerto Colombia - Atlántico.

Icons indicate efficiency and richness ranks for each order and each sampling method.

Order	Total Richness	Pitfall	Malaise	Fogging
 Hymenoptera	137	29 	102 	25 
 Coleoptera	125	47 	77 	25 
 Diptera	87	9 	80 	2 
 Hemiptera	36	7 	18 	14 
 Psocoptera	9	- 	2 	9 
 Blattodea	4	4 	1 	- 
 Mantoidea	5	1 	1 	3 
 Orthoptera	5	3 	2 	- 
 Collembola	2	2 	- 	- 
 Embioptera	1	1 	- 	- 
 Neuroptera	1	- 	1 	- 
 Zygentoma	1	1 	- 	- 

	Species count	Sample Coverage	Total richness
	> 10	> 80 %	> 50 %
	≥ 4	> 50 %	> 30 %
	< 3	occasional sample	
	No reports		

Figure 2

Distribution of abundance by each order of insects in each of the collection methods evaluated

Each bar indicates the number of morphospecies (observed richness) for each order.

Different colors in each order indicate richness by type of trap. Icons are meant as a proxy of a generic representative per order.

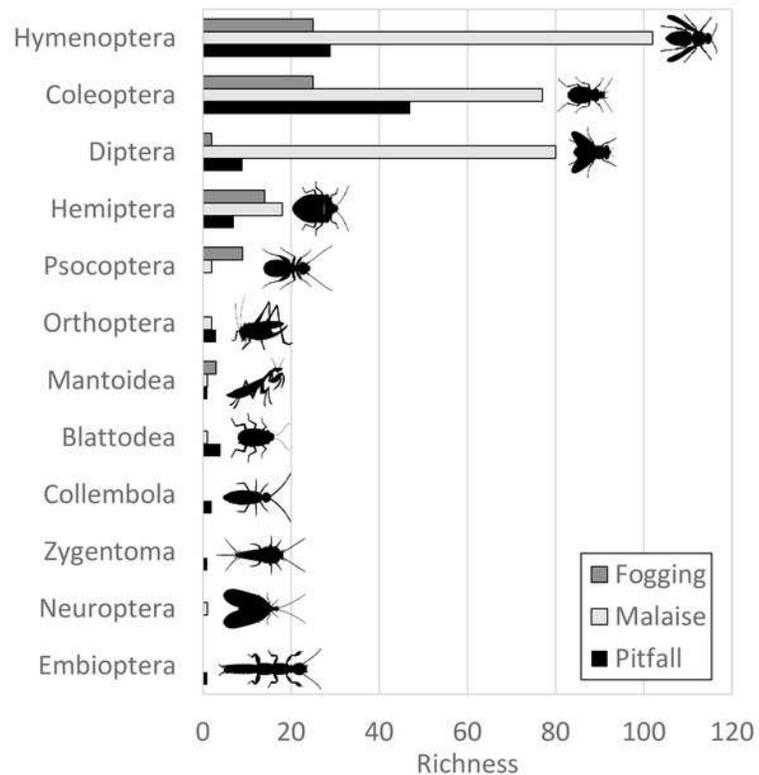


Figure 3

Percentage of sampling coverage for each collection method

Each line represents the dynamic behavior of the richness relative to the total sampling. Continuous line represents rarefaction, while dotted line represents extrapolation. Symbol indicates estimated richness. All three types of traps have an estimated coverage larger than 90% overall. Individual orders have different behavior (Supplemental Figure 1)

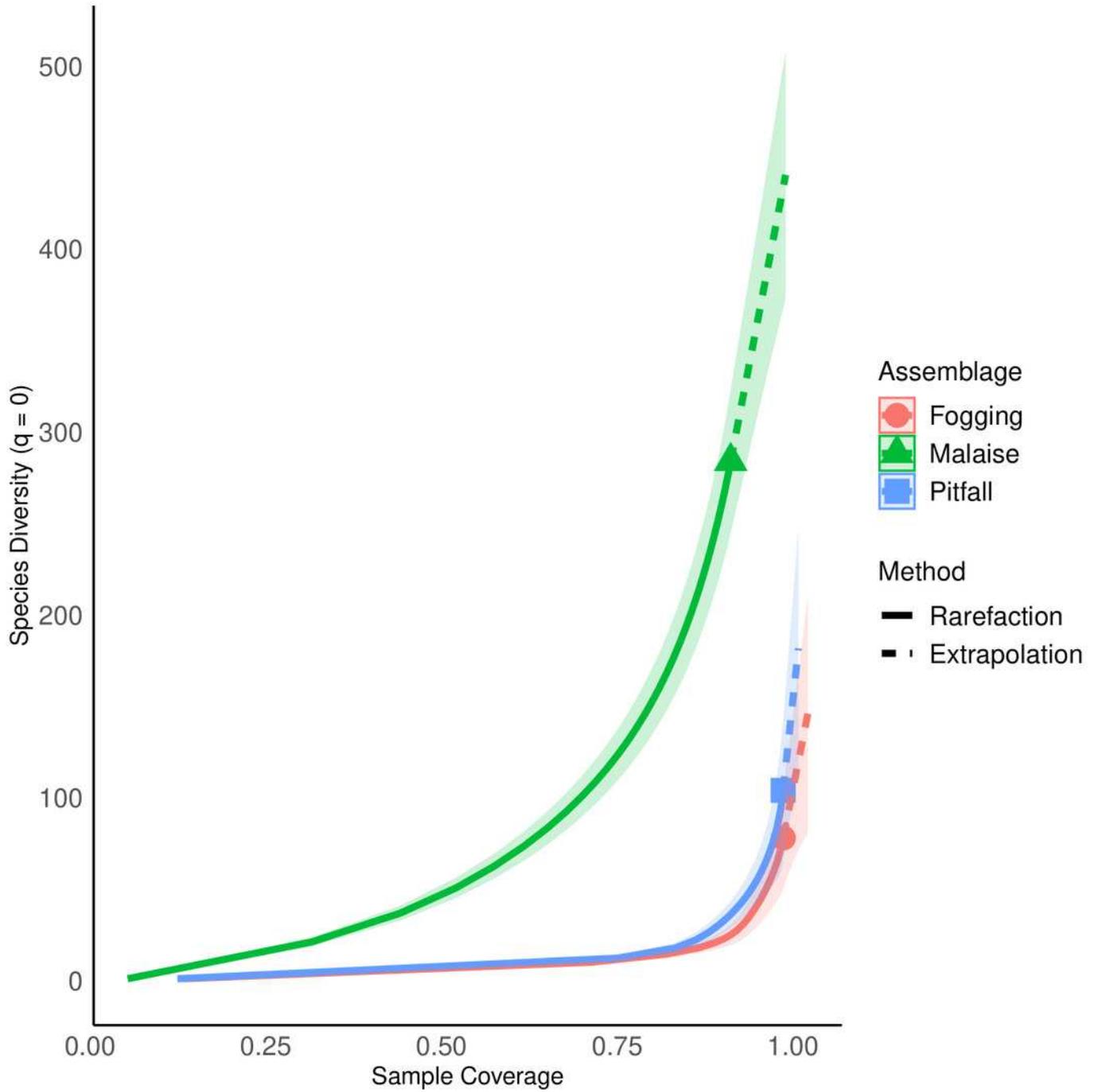


Figure 4

Rarefaction and extrapolation of Diversity based on Hill numbers for Malaise, Fogging and Pitfall sampling methods

(A) Species richness ($q=0$), (B) Shannon Diversity, $q = 1$, (C) Simpson Diversity $q = 2$. Each curve is computed from 999 bootstraps. Continuous line indicates rarefaction. Dashed line indicates extrapolation. Symbol indicates estimated richness for each index. Shaded area represents 95% confidence intervals.

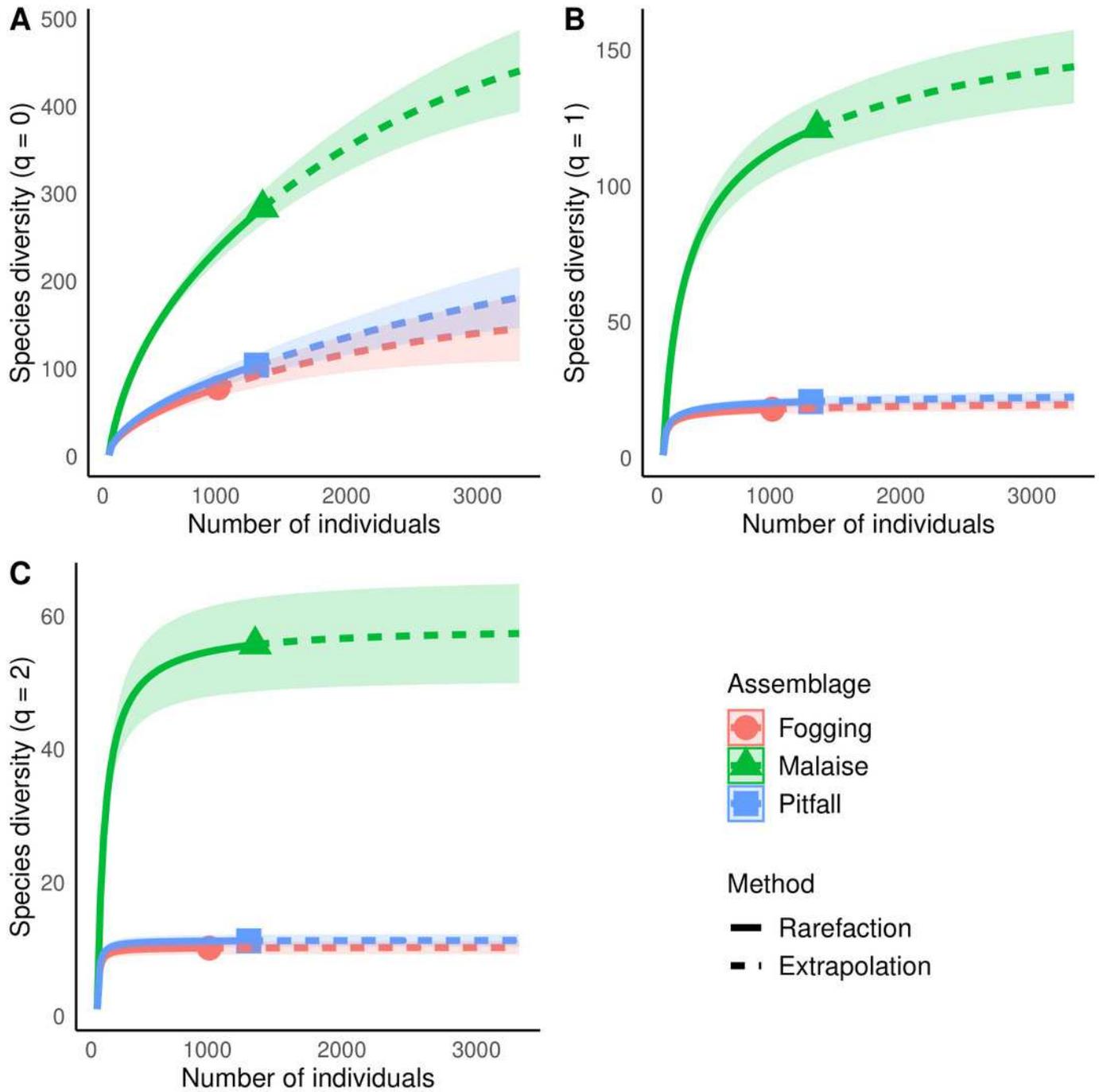


Figure 5

Beta diversity of insect communities with three collection methods evaluated in a relict of Dry Forest in Puerto Colombia

(A-C) Venn diagrams of shared (A) morphospecies, (B) families and (B) orders among methods, respectively. Darker shades of gray indicate higher counts. (D) Partition of the diversity of β SOR (blue dashed line) between β SIM (black dotted line) and β SNE (red continuous line). (E). Clustering using the average of the β SIM dissimilarity of the species composition.

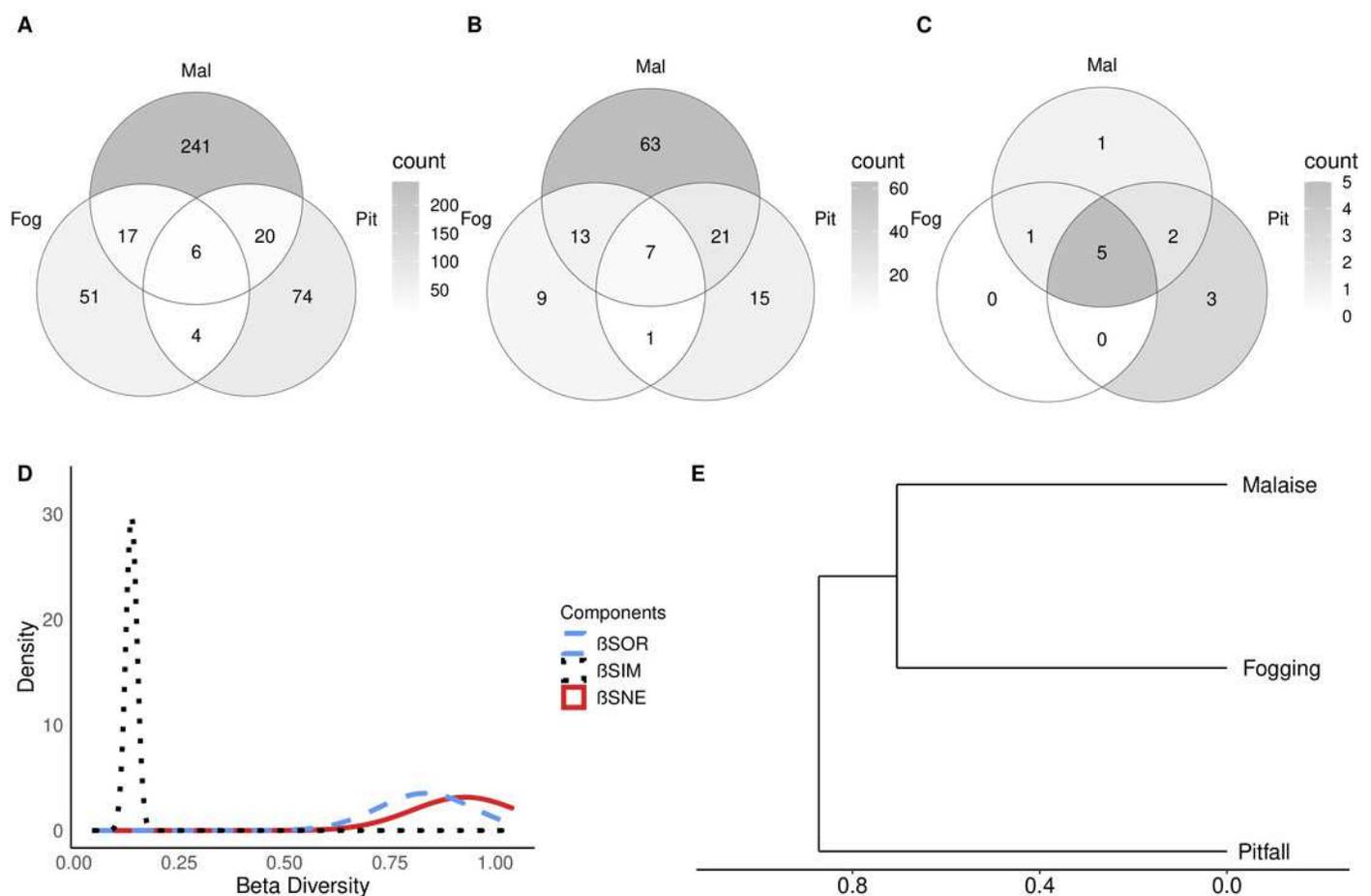


Figure 6

Distribution of richness by detected guilds and collection methods

Each bar represents the number of morphospecies belonging to a particular guild. Different colors represent different traps.

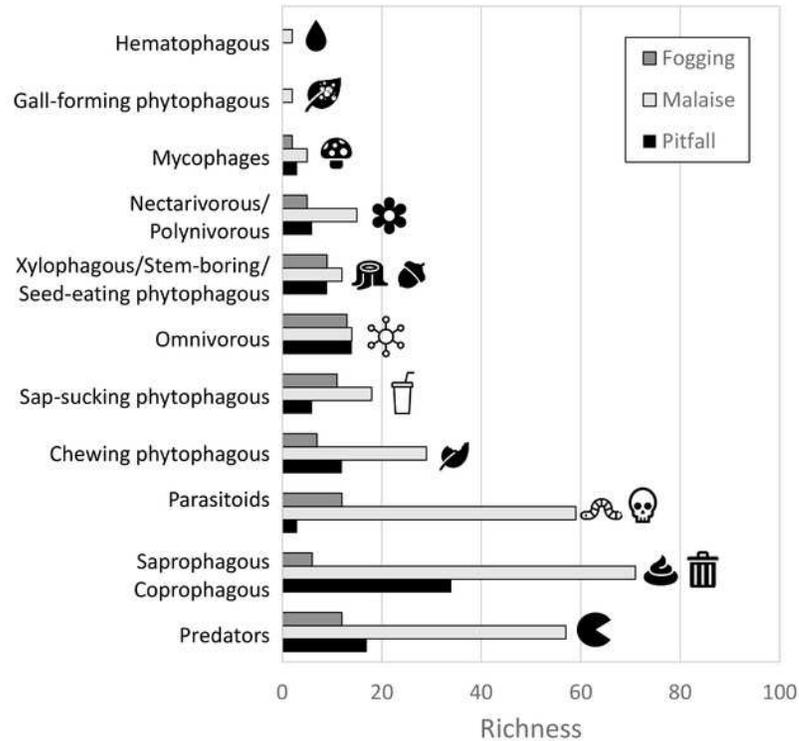


Figure 7

Richness by trophic guilds and efficiency of three different collection methods and their efficiency in a relict of dry forest in the municipality of Puerto Colombia - Atlántico.

Icons indicate efficiency and richness ranks for each guild and each sampling method.

Order	Total Richness	Pitfall	Malaise	Fogging
 Saprophagous/Coprotophagous	100	6 	71 	34 
 Predators	79	12 	57 	17 
 Parasitoids	73	12 	59 	3 
 Chewing phytophagous	43	7 	29 	12 
 Sap-sucking phytophagous	32	11 	18 	6 
 Omnivorous	31	13 	14 	14 
 Xylophagous/Stem-boring	22	9 	12 	9 
 Seed-eating phytophagous				
 Nectarivorous/Polynivorous	19	5 	15 	6 
 Mycophages	9	2 	5 	3 
 Gall-forming phytophagous	2		2 	
 Hematophagous	2		2 	

	Species count	Sample Coverage	Total richness
	> 10	> 80 %	> 50 %
	≥ 4	> 50 %	> 30 %
	< 3	occasional sample	
	No reports		

Table 1 (on next page)

Richness of insect trophic guilds and their equitativity

Richness and equity index J for three different collection methods, as a proxy for functional diversity distribution across methods.

1 **Table 3. Richness of insect trophic guilds with three different collection methods in a**
2 **relict of dry forest in the municipality of Puerto Colombia – Atlántico.**
3

	Pitfall	Malaise	Fogging
Trophic guilds present	9	11	9
Equitativity J	0.89	0.84	0.95

4