

Assessing the Climatic Vulnerability of *Micrurus* *sangilensis* (Niceforo Maria, 1942) under Future Scenarios

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1 **Assessing the Climatic Vulnerability of the *Micrurus sangilensis* (Niceforo Maria, 1942)**
2 **under Future Scenarios**

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19 **Abstract.** The vulnerable *Micrurus sangilensis* commonly known as the Santander coral snake
20 distributes in dry and montane forests, ecosystems under severe anthropogenic pressure in
21 northeastern Colombia. The habitat of this serpent is fragmented, and climate change may
22 further intensify risks to the vegetation structure. We assessed whether the current distribution
23 of the snake may be altered under different scenarios with climate change in the 2040-2060
24 years; aiming to recognize conservation priority areas. With ecological niche modeling we
25 calculated current values of stability in the distribution range of the species, for the most
26 conservative emission scenarios of Socio-Economic Pathways (SSP) 126, and 245; and the
27 expected greater emissions 585 within five different global circulation models. We also
28 escalated an index of vulnerability to land use change to 2050 in the remaining areas for the
29 species, detecting prioritizing conservation zones. Our findings reveal a nearly 25% consistency
30 of loss in the three SSP scenarios, while gaining stability varies between different GCMs. Over
31 37% of remaining suitable areas were categorized as highly vulnerable to land-use change,
32 especially at elevations between 900 and 2000 m. We emphasize the need to integrate *M.*
33 *sangilensis* habitats into Colombia's protected area network, restore degraded ecosystems, and
34 establish ecological corridors to mitigate fragmentation. While the most vulnerable to changing
35 areas appear to be the ones with critical requirements for conservation; We recommend
36 targeting conservation efforts in areas of low to medium vulnerability to change, which are less
37 likely to undergo significant modifications over the next 30–40 years

38

39 **Keywords.** Biodiversity conservation, climate change, ecological niche modeling, fragmented
40 landscapes, *Micrurus sangilensis*

41

43 The rapid changing patterns in climatic regimes have become a central concern in
44 conservation sciences (Lovejoy, 2006; Young et al., 2011; Yu et al., 2014; Upadhyay, 2020;
45 Fuentes et al., 2023). In the Americas, home to multiple biodiversity hotspots, ecosystems are
46 increasingly at risk as altered climatic conditions force species to shift their historical
47 distribution ranges toward new latitudinal and altitudinal zones in search of suitable habitat
48 conditions (Allentoft and O'Brien, 2010; Bellard et al., 2012; Vicenzi et al., 2017; Archis et al.,
49 2018). Furthermore, range-restricted and threatened species often lack the dispersal capacity to
50 cope with these changes across fragmented landscapes with rapid shifting, among others,
51 vegetation structures (Kwak and Freeman, 2010; Bestion et al., 2015; Upadhyay, 2020; Fuentes
52 et al., 2023). While climate change is undoubtedly a major threat to already vulnerable
53 ecosystems, rapid and drastic land-use changes may pose an even more immediate challenge,
54 this compromises the structural integrity of habitats and increases barriers to species movement
55 (Bellard et al., 2012; Kittel, 2013; Staudt, 2013; Bush et al., 2017; Cobos et al., 2018). These
56 combined pressures underscore the urgent need for conservation strategies that enable the most
57 threatened, range-restricted species to respond to both climate and land-use changes,
58 particularly in fragmented landscapes (Heller and Zavaleta, 2009; Mawdsley et al., 2009; Rands
59 et al., 2010).

60 In South America Andean ecosystems are important reservoirs of rich and complex
61 biodiversity (Sarkar et al., 2009; Ramirez-Villegas et al., 2014; Bax and Francesconi, 2019),
62 where inhabiting hundreds of endemic and range restricted species, some specialized to the
63 typical habitats in ranges and associated environments; however, also among the most
64 threatened ecosystems in the world (Young et al., 2011; Bax and Francesconi, 2019; Noh et al.,
65 2020). Among many other groups affected by the changing landscape conditions in the Andes,
66 reptiles exhibit high sensitivity to climate change and habitat loss, mainly in relation to their

67 reproductive processes (Huey et al., 2009; Gamble, 2010). However, they should be given
68 higher priority, particularly in the context of climate change impacts (Gumbs et al., 2018).
69 Residing in delicate ecosystems already imperiled by deforestation and habitat degradation,
70 many reptile species are categorized as threatened according to the IUCN Red List (Arredondo
71 et al., 2015; Bolívar et al., 2016; Páez et al., 2016; Calderón et al., 2019; Hladki et al., 2019;
72 Rainwater et al., 2022). They confront altered temperature and precipitation patterns, imperiling
73 their ecological niches and reproductive cycles (Brown and Shine, 2006; Gamble, 2010). The
74 ramifications extend to the intricate balance of their habitats, where some with restricted ranges
75 and specialized ecological requirements are particularly susceptible to environmental
76 perturbations (Holt, 1990; Moraes and Recchia, 2011).

77 Among reptiles, the family Elapidae, which includes coral snakes (*Micrurus*) have
78 species with wide distributions such as (*Micrurus dumerilli* and *M. dissolcucus*), and a high
79 tolerance for climatic regimes, as *M. mipartitus* inhabiting from 0 to near 2500 m (Rey-Suárez
80 et al., 2016; Herrera-Lopera et al., 2018; Pitalua et al., 2018; Río-Soto et al., 2018). However,
81 there are also some snakes with reduced distribution such as *M. sangilensis*, which is especially
82 affected by the modification, degradation, and loss of natural habitat (Hladki et al., 2019; Flórez
83 and Montoya-Cruz, 2023).

84 *Micrurus sangilensis*, commonly known as the Santander coral snake, is a triad-colored
85 species distinguished by specific ring patterns and the absence of supracloacal keels. It differs
86 from its close relatives, *M. dissolcucus* and *M. dumerilli*, by having 16–22 triads and a typical
87 length of around 60 cm (Morales-Betancourt et al., 2015) (Fig. 1). Endemic to Colombia, its
88 narrowed distribution within the departments of Cundinamarca, Boyacá, and Santander (Roze,
89 1996; Campbell et al., 2004; Caicedo-Portilla and Lynch, 2015) and more recently detected in
90 Casanare (Flórez and Montoya-Cruz, 2023). This species limited to moderate elevations (800–
91 2800 above sea level) along the Middle Magdalena River Basin, primarily inhabiting the

92 vulnerable dry forests (Campbell et al., 2004). Currently classified as Vulnerable by the IUCN,
93 its historical records show an estimated extent of occupation near 12000 km² (Caicedo-Portilla
94 and Lynch, 2015; Hladki et al., 2019). Despite its conservation status, critical knowledge gaps
95 remain regarding its population size, reproductive dynamics, and habitat requirements, which
96 are essential knowledge when working for effective conservation planning (Van Teeffelen et
97 al., 2012; Caicedo-Portilla and Lynch, 2015). Species like *M. sangilensis* may face acute risks
98 not only from climate change but also from rapid structural changes in vegetation due to habitat
99 degradation (Caicedo-Portilla and Lynch, 2015; Hladki et al., 2019; Flórez and Montoya-Cruz,
100 2023).

101 Given the ongoing changes in land use and the potentially altered climatic regimes in
102 rainfall and temperature within the degraded dry forests, typical habitat of the restricted
103 distribution of *M. sangilensis*, it is necessary to develop management strategies predicting the
104 potential shifts in the species' distribution under future climate scenarios. This research aims to
105 assess the species' potential distribution under different Shared Socioeconomic Pathways (SSP)
106 under climate change, through a vulnerability to change index for land use, using Ecological
107 Niche Modeling Methodologies (ENM) which are among the most widely used and effective
108 tools for comparing and predicting historical and future scenarios (Peterson et al., 2011; Rangel
109 and Loyola, 2012). By relating environmental and climatic variables with species occurrence
110 data, ENM offers insights into species' range dynamics (Alvarado-Serrano and Knowles, 2014;
111 Ramirez-Villegas et al., 2014; Mota-Vargas and Rojas-Soto, 2016; Moreno-Contreras et al.,
112 2020; Sales et al., 2020). This assessment serves as a crucial resource for conservation planning
113 strategies, highlighting areas for immediate and future protection efforts in landscapes
114 increasingly fragmented by agriculture and urbanization.

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117

MATERIAL AND METHODS

118

119 *Occurrences and accessibility area (M)*

120 We obtained records of *M. sangilensis* from the Global Biodiversity Information
121 Facility database (GBIF <https://doi.org/10.15468/dl.9cjqyy>) and literature (Flórez and
122 Montoya-Cruz, 2023), in total 59 Records were used for analyses after deleting duplicates and
123 reviewing each locality consistency to the known species historical range, following the
124 methodology based on Cobos and Bosch, (2018). The accessible area, commonly named as M
125 (Soberon and Peterson, 2005), represents the geographic region that the species could have
126 potentially occupied over relevant evolutionary time frames, based on its dispersal ability and
127 known biogeographic barriers, this M is the spatial extent from which environmental variables
128 are cropped to calibrate the model and generate predictions; by considering at least one
129 occurrence in terrestrial ecoregions from The Nature Conservancy (Dinerstein et al., 2017),
130 using the ArcMap 10.8 software (Fig. 2).

131

132 *Environmental variables*

133 We obtained climatic variables to characterize the climatic niche of the *M. sangilensis*
134 from the WorldClim 2.1 database (Fick and Hijmans, 2017) at a 30 arc-second resolution (~1
135 km²). We exclusively used 15 biovariables, excluding bio 8 (mean temperature of wettest
136 quarter), bio 9 (mean temperature of driest quarter), bio 18 (precipitation of warmest quarter),
137 and bio 19 (precipitation of coldest quarter); these exclusions reduce redundancy and avoid
138 potential collinearity, overfitting or misrepresenting the species' niche model (Escobar et al.,
139 2014; Table 1 in supplementary material). Variables were masked and cropped to the extension
140 of the accessible area of *M. sangilensis*. Because some highly correlated variables can be
141 important for the biology of the species, we created different sets from the 15 biovariables for

142 niche modeling evaluation (Cobos and Bosch, 2018; Echeverry-Cárdenas et al., 2021). The
143 specific sets are shown in Table 2 in supplementary material.

144

145 *Ecological niche modeling (ENM)*

146 We randomly allocated 80% of the occurrence records for model training and the
147 remaining 20% for evaluation, using the Maxent algorithm (Elith et al., 2011) implemented
148 through the kuenm package in R (Cobos et al., 2019). We assessed multiple levels of model
149 complexity following the approach of Warren and Seifert (2011) by varying the regularization
150 multiplier from 0.1 to 1 in increments of 0.1, and then testing values of 2, 3, 4, and 5. Model
151 selection was based on performance metrics provided by Kuenm, including the AUC ratio,
152 omission rate, and AIC. The AUC (Area Under the Curve) ratio measures the model's ability
153 to discriminate suitable versus unsuitable areas, while the AICc (corrected Akaike Information
154 Criterion) evaluates model parsimony by balancing goodness-of-fit and complexity (Phillips et
155 al. 2006; Pearson et al. 2007; Warren and Seifert 2011; Arango-Lozano et al., 2025). The
156 selected model was projected without applying extrapolation modes (Extrapolation or
157 Clamping in Maxent algorithm) onto the accessible area under future climate scenarios,
158 specifically Shared Socioeconomic Pathways (SSPs) representing minimal and moderate
159 climatic changes (SSP126 and SSP245), as well as the most extreme scenario (SSP585), a high-
160 emissions scenario associated with continued fossil fuel use and the most severe climate change
161 (Echeverry-Cárdenas et al., 2021; Arango-Lozano et al., 2025) for the cumulative years 2041–
162 2060.

163 We used five Global Circulation Models (GCMs) for comparing results as: HadGEM3-
164 GC31-LL (GCM1), IPSL-CM6A-LR (GCM2), ACCESS-CM2 (GCM3), EC-Earth3-Veg
165 (GCM4) and UKESM1-0-LL (GCM5). Each GCM was chosen for its unique approach in
166 simulating climate dynamics, using distinct datasets, including land cover, oceanic circulation,

167 and atmospheric processes (Yu et al., 2014; Padhiary et al., 2020). The variety of GCMs
168 enhances model robustness, enabling us to assess areas of agreement and divergence via a
169 broader perspective on future climate variability (Reshmidevi et al., 2018; Padhiary et al.,
170 2020). For example, HadGEM3-GC31-LL focuses on ocean-atmosphere interactions, while
171 UKESM1-0-LL includes comprehensive land-use feedback (Reshmidevi et al., 2018; García-
172 Franco et al., 2020). We reclassified models resulted in presence/absence maps using a
173 consistent threshold across models (10 percentile training presence Cloghold in Maxent
174 algorithm).

175

176 *Gain, loss, and stability*

177 We evaluated the current and future potential distribution by analyzing pixel counts (and
178 then its scaled values to Km²). First, we calculated the percentage of pixels gained, lost, and
179 stabilized in each scenario (SSP and GCM) relative to the current distribution. We then
180 summarized the gained and stabilized pixels for each SSP, identifying areas that remained stable
181 across future GCMs. To pinpoint regions of consistent stability, we focused on pixels that
182 persisted in four or more GCMs, allowing us to observe the spatial assemblage of stable areas
183 under future conditions across all five GCMs.

184

185 *Vulnerability to change*

186 Finally, to detect any potential vulnerable areas of habitat lost in the species potential
187 distribution, we superpose the current and future (assembly pixels in four or more GCMs)
188 ranges with a “Vulnerability of land cover to anthropogenic change raster” (Esri, 2024). This
189 layer shows areas where natural vegetation such as forest could be converted to agriculture and
190 urban lands by 2050, based on the predictions of human-induced land changes. Further, the
191 layer includes regions susceptible to expansion of agricultural and urban footprints, excluding

192 forecasts for unchanged land cover types like forests unless they are converted to agriculture or
193 urban areas (available data in: <https://livingatlas.arcgis.com/landcover-2050/>).
194 The vulnerability data ranges from zero to one, indicating varying degrees of susceptibility to
195 natural cover transformation. We classified the raster values into three distinct categories: low
196 vulnerability (0.0 - 0.3), moderate vulnerability (0.3 - 0.7), and high vulnerability (0.7 - 1) for
197 the species. Areas with high vulnerability values (0.7 - 1) are identified as the most critical
198 zones for this snake. These areas are also deemed most at risk of change in future climate
199 scenarios (Arango-Lozano et al., 2025).

200

201 RESULTS

202 *Ecological niche model selection*

203 Out of 210 evaluated models, only 3 fulfilled the kuenm criteria, and to streamline niche
204 model comparisons between scenarios, we selected the first exhibiting the lowest AIC and AUC
205 results M_0.7_F_lq_Set 5. Set 5 consists of biovariables 2, 3, 4, 7, 16, and 17, representing a
206 combination of precipitation and temperature conditions. (Table 2, Table 3 in supplementary
207 material).

208

209 *Gain, loss, and stability*

210 For the current scenario of *M. sangilensis*, 19873 predicted pixels were identified, which
211 translate into over 16879 km² of suitable environmental conditions (Fig. 4A). Building on this
212 baseline, future projections indicate substantial variability in potential range across General
213 Circulation Models (GCMs) and Shared Socioeconomic Pathways (SSPs), yet certain
214 consistent patterns emerge that strengthen confidence in key findings. Across all scenarios,
215 significant stability areas are consistently predicted, particularly in the lower-emission scenario

216 (SSP126). For instance, stability remains high in models M2 and M4 under SSP126, with over
217 80% of predicted pixels showing little change (Fig. 3).

218 As emissions concentrations increase (SSP245 and SSP585), a marked rise in habitat
219 gain is observed. Under SSP585, models such as M1 and M5 predict substantial habitat gains,
220 with M5 showing a gain of 38.7% of pixels. This indicates possible new areas as climatic
221 conditions shift, though this expansion comes with increased variability in predicted losses,
222 particularly in M3 and M4 (Fig. 3). The appearance of these gained pixels under higher
223 emissions scenarios reflects potential new suitable habitats as the climate changes. While the
224 model variability shows the inherent uncertainties tied to different GCMs, the overall trends of
225 habitat stability and gains under higher emissions scenarios are robust across models.

226

227 *Vulnerability to change*

228 We identified a mean reduction of at least 25% of suitable future areas for the species
229 with respect to the current scenario. However, when contrasting the stability values with the
230 vulnerability to change 2050 layer, we recognized that at least 37% of the remnant areas in the
231 different future scenarios (each SPP) may be experiencing a high vulnerability, 30% a medium
232 and almost just 23% a low vulnerability (Fig. 4). There was identified loss of areas in all
233 elevation between 900 and 2000 m, Furthermore, these areas consistently show both lost in
234 predicted presence and where not, are the ones with greater vulnerability values (0.7 - 1).

235

236 DISCUSSION

237 The predictions for *Micrurus sangilensis* are influenced by the inherent uncertainties
238 associated with the different General Circulation Models (GCMs), each of which makes varying
239 assumptions about climate processes (Reshmidevi et al., 2018; Padhiary et al., 2020). By
240 narrowing our analysis to pixels that remained stable across four or more GCMs, we were able

241 to highlight regions that are likely to maintain suitable conditions for *M. sangilensis* even under
242 varying future climate scenarios. This method provided a clearer spatial picture of potential
243 refugia and stable areas, ensuring that our findings reflect consistent patterns of stability rather
244 than relying on any single model. The identification of these stable regions across multiple
245 GCMs strengthens the forecasting future conditions for this coral snake.

246 Our study projects a significant contraction in the potential suitable habitat for *Micrurus*
247 *sangilensis* due to climate change across all SSP scenarios. These findings are consistent with
248 other studies on coral snakes, such as *M. lemniscatus*, where a similar vulnerability to habitat
249 loss in lowland regions was observed (Terribile et al., 2018). For *M. fulvius*, a potential range
250 shift into unsuitable areas was predicted, highlighting the species' inability to keep pace with
251 changing climatic conditions (Archis et al., 2018). Similarly, *M. brasiliensis* faces the potential
252 loss of 60% of its ideal habitat under future climate projections (Caten et al., 2017), further
253 underscoring the widespread impact of climate change on coral snake distributions.

254 In the case of our study species *M. sangilensis*, our results indicate a potential loss of
255 25% of its ideal habitat by 2040–2060, with the species currently occupying degraded dry forest
256 ecosystems, some of the most threatened habitats globally (Miles et al., 2006). Additionally, a
257 30% increase in the vulnerability of remaining suitable areas, particularly from land-use
258 changes in forests (Galindo-Cruz et al., 2024). Within less than two decades, these critical
259 habitats could undergo significant alterations, threatening the species' long-term survival
260 (Hladki et al., 2019).

261 The broader implications of these results become clear when considering that future
262 suitable habitats for this species may be out of reach. As observed in other species, the pace of
263 climate-driven habitat change may exceed the distribution capabilities of an animal (Schloss et
264 al., 2012; Bush and Hoskins, 2017; Sales et al., 2020). Furthermore, factors such as habitat
265 fragmentation and population isolation could further limit the species' ability to migrate to

266 newly suitable areas (Le Galliard et al., 2012; Bestion et al., 2015). As a result, even with new
267 habitats emerging and stabilizing under future climatic conditions, the species may not be able
268 to reach or colonize them effectively (Loarie et al., 2009; Le Galliard et al., 2012; Sales et al.,
269 2020; Arango-Lozano et al., 2025).

270 The conservation implications of our results underscore the need for both immediate
271 and long-term strategies. Areas of high vulnerability (with values between 0.7 and 1), as
272 identified in this study, represent critical zones where habitat transformation is likely imminent.
273 However, it is essential to recognize that focusing solely on these high-risk areas may not
274 prevent transformation, as rapid environmental changes might already be underway (Global
275 Forest Review, 2024). This introduces a crucial discussion point: should conservation efforts
276 be prioritized in areas of highest vulnerability, or should a broader approach be taken, including
277 areas with moderate vulnerability?

278 Given the lack of clear protocols for prioritizing conservation actions in snakes
279 (Terribile et al., 2009; Andrade-Díaz et al., 2019), it might be strategic to include areas across
280 the vulnerability spectrum. For instance, regions with moderate vulnerability ('yellow zones')
281 are less likely to undergo immediate change compared to high-risk areas ('red zones'), but they
282 remain vital for habitat connectivity and the long-term survival of species like *Micrurus*
283 *sangilensis*. Conservation efforts in these areas could help buffer against habitat fragmentation
284 and provide refugia as the more vulnerable areas degrade.

285 In Colombia, existing conservation plans and frameworks provide an opportunity to
286 safeguard the critical habitats for *Micrurus sangilensis*. The dry forests of Santander, where
287 suitable areas for this species have been identified, are adjacent to over 60 different protected
288 areas (Fig. 5). These areas include National Natural Parks, regional integrated management
289 districts, national reserves, and forest reserves, all recognized under the National Register of

290 Protected Areas (RUNAP). This network of protected zones represents a significant collective
291 effort to preserve the country's biodiversity.

292 Although *M. sangilensis* may benefit from the protection offered by some of these
293 existing areas, additional conservation strategies will be vital (Mi et al., 2023). Expanding the
294 coverage of protected areas, restoring degraded habitats, and establishing ecological corridors
295 to link fragmented landscapes are key steps to enhancing the resilience of this species under
296 changing climate conditions (Terribile et al., 2009; Andrade-Díaz et al., 2019). Moreover,
297 incorporating climate-adaptive conservation actions into these plans could help ensure that *M.*
298 *sangilensis* can persist in its shrinking habitat. Ongoing monitoring of land-use changes and the
299 potential threats posed by human activities will also be essential in aligning conservation efforts
300 with the species' emerging vulnerabilities (Fordham et al., 2012; Mi et al., 2023). A
301 collaborative, multidisciplinary approach is crucial to create effective conservation strategies,
302 safeguard the species' habitat, and ensure the long-term survival of the Santander coral snake.

303

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308

309

310 SUPPLEMENTARY MATERIAL

311 The supplementary materials include three (3) tables describing the use of climatic variables
312 and the results selected in this study ecological niche models, annexed to this manuscript.
313 Additionally, we have made available online the resulted raster files with current and projected
314 future species distributions at: OSF <https://osf.io/h9srj/>.

315

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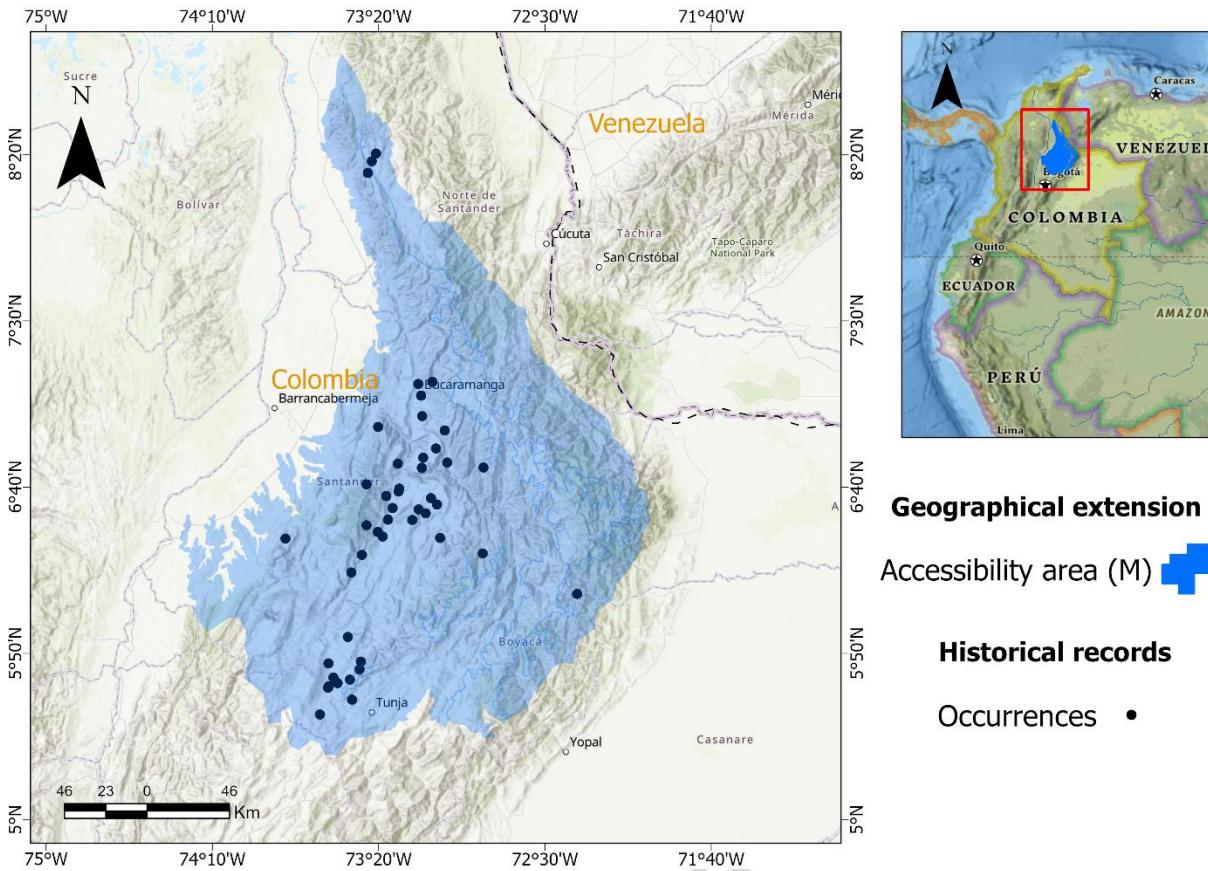


528

529 **Fig. 1.** Photograph of a living adult of *Micrurus sangilensis* with its characteristic coral pattern.

530 Photo by: Elson Meneses-Pelayo.

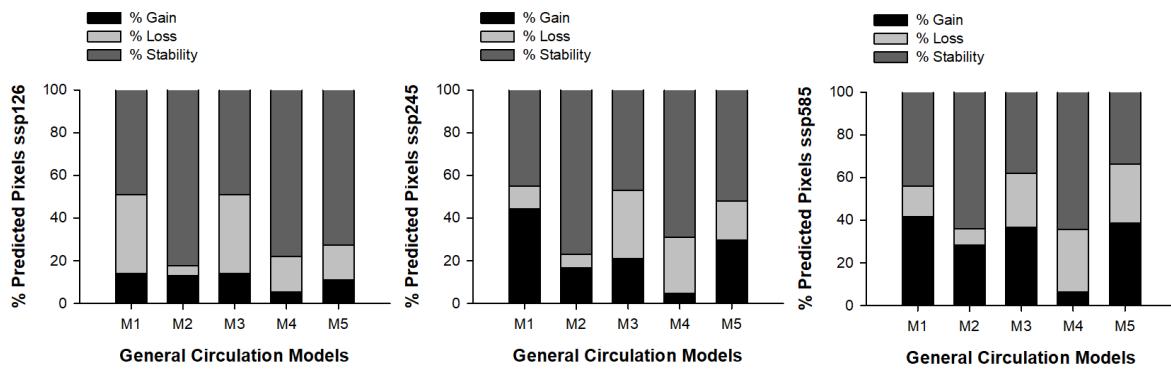
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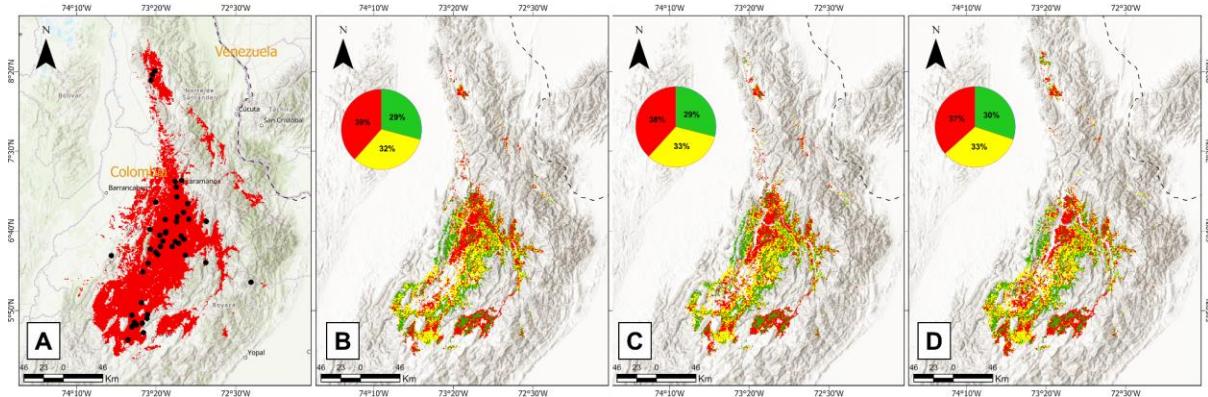
533 **Fig. 2.** Historically known distribution of *Micrurus sangilensis* and its generated accessible area
534 (M).

535



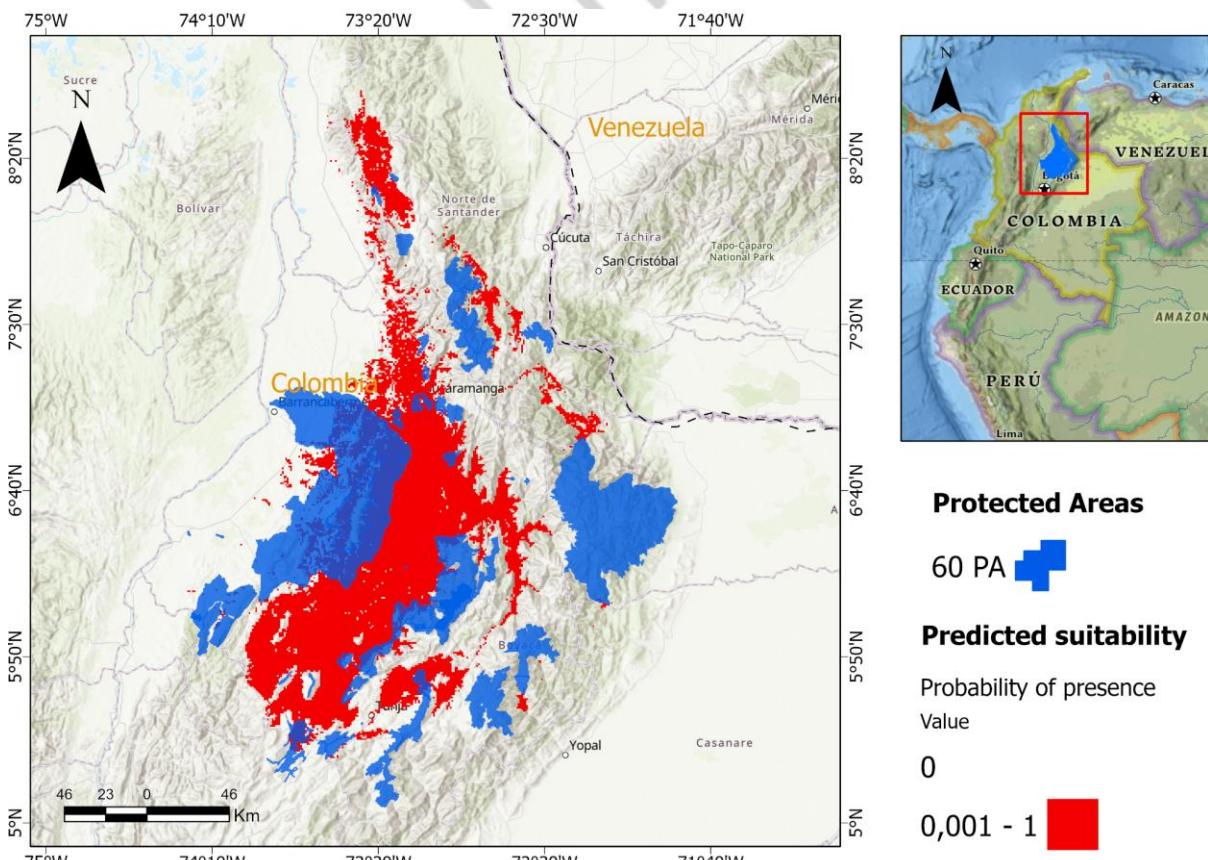
536

537 **Fig. 3.** Percentage of gained loss and stability of predicted pixels (occurrence likelihood = 1)
538 under future scenarios for *Micrurus sangilensis*. Results are shown for different General
539 Circulation Models (GCMs, labeled M1–M5) across the Shared Socioeconomic Pathways
540 (SSPs: ssp126, ssp245, ssp585).



542 **Fig. 4.** Comparison of suitable conditions for *Micrurus sangilensis* across different scenarios.
 543 The subsequent maps depict the accessibility area (M) under various shared socioeconomic
 544 pathways and timeframes: (A) current conditions with occurrences (black dots), (B) assembly
 545 of consistent ideal areas in SSP126, (C) assembly of consistent ideal areas in SSP245, (D)
 546 assembly of consistent ideal areas in SSP585. Colored pixels indicating values of vulnerability
 547 to change in the land use: green = Low, yellow = Mid, red = High.

548



550 **Fig. 5.** Description of the sloping between the suitable areas for the distribution of *Micrurus*
551 *sangilensis* and surrounding protected areas in the surrounding region of Santander Colombia.
552