

# Habitat transformation reshapes diversity and community structure of amphibians and reptiles in the Eastern Andes

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## Abstract

Land-use change is a major driver of biodiversity loss in tropical montane ecosystems, yet its effects on herpetofaunal community structure remain insufficiently understood at local scales. We evaluated patterns of diversity and community structure of amphibians and reptiles across a gradient of habitat transformation in southwestern Cundinamarca, Colombia. Field surveys conducted between 2015 and 2022 using visual encounter surveys recorded 57 species and 608 individuals. Forest habitats supported the highest species richness and diversity, whereas coffee crops and open areas exhibited reduced richness and simplified assemblages. Rank–abundance patterns revealed greater dominance in open habitats and higher evenness in forests, consistent with environmental filtering processes. Amphibians were more strongly associated with forest environments, whereas reptiles showed broader habitat tolerance and higher representation in disturbed habitats. In addition, we compiled a comprehensive regional inventory based on field data, biological collections, and biodiversity databases, documenting 86 species of herpetofauna. Overall, our results show patterns consistent with environmental filtering, where habitat transformation is associated with increased dominance of generalist species and simplified community structure. These findings highlight the importance of conserving forest remnants and maintaining heterogeneous landscapes to support herpetofaunal diversity in the Colombian Andes.

## Key Words

agroecosystems, anthropic disturbance, biodiversity conservation, habitat use, herpetofauna inventory

## Introduction

In Colombia, Andean ecosystems have undergone extensive transformation, particularly in Andean and sub-Andean forests, which concentrate the highest levels of productive activity in the country and, consequently,

strong anthropogenic pressures mediated by ranching, agriculture, infrastructure development and mining (Correa-Ayram et al. 2020; Rubiano et al. 2026). The Andean region harbors approximately 53% of the country's total herpetofauna (Romero et al. 2009), including a high number of endemic species (Bernal and Lynch 2008; Armesto

and Señaris 2017). At a broader scale, the tropical Andes are recognized as a global biodiversity hotspot (Myers et al. 2000; Vasconcelos et al. 2019), characterized by exceptional levels of endemism and a high concentration of threatened species (Tobar-Suárez et al. 2022; IUCN 2024; Vásquez-Restrepo and García-Cobos 2026). Additionally, a substantial proportion of these species occurs outside protected areas, increasing their vulnerability to ongoing land-use change, particularly in human-dominated landscapes (Nori et al. 2015; Bax and Francesconi 2019; Vásquez-Restrepo and García-Cobos 2026).

Two main processes have been proposed to explain how ecological communities are structured, both of which are relevant to understanding the consequences of anthropogenic pressures. These pressures affect not only species richness but also patterns of dominance, evenness, and species' interactions (Ernst et al. 2006; Gardner et al. 2007). Limiting similarity hypothesis suggests that species with high niche overlap experience competitive exclusion, leading to reduced overall abundance but favoring the persistence of species that exploit different resources (MacArthur and Levins 1967; Córdova-Tapia and Zambrano 2015). In contrast, environmental filtering emphasizes how abiotic conditions allow only species with certain traits to persist, thereby shaping both richness and abundance (Brüning et al. 2018; Zabala-Forero and Urbina-Cardona 2021; López-Bedoya et al. 2022). In herpetofaunal assemblages, habitat transformation is frequently associated with environmental filtering due to structural changes, edge effects, and altered resource availability (Granda-Rodríguez et al. 2025; Palomino-Cuéllar and Urbina-Cardona 2025).

Amphibian and reptile assemblages in tropical landscapes are strongly affected by land-use changes linked to agriculture, livestock, and urban expansion (Ghosh and Basu 2020; Iglesias-Carrasco et al. 2023). These activities drive habitat loss, degradation, and fragmentation, producing heterogeneous landscapes composed of remnant forest patches embedded within human-modified matrices (Thompson et al. 2015; Cervantes-López and Morante-Filho 2024). Disturbed habitats are therefore often characterized by simplified assemblages dominated by a few generalist species. Responses to habitat transformation are expected to differ between amphibians and reptiles due to variation in physiological and ecological traits. Amphibians, with their permeable skin, dependence on moisture for reproduction, and narrow thermal tolerances, are generally more sensitive to environmental changes (Wanger et al. 2010; Palacios et al. 2013; Méndez-Narváez 2014; Sankararaman and Miller 2024). Reptiles, by contrast, typically exhibit broader tolerance to thermal and hydric variation, which enables them to persist (or even increase in abundance) in disturbed environments (Thompson et al. 2015; Cordier et al. 2021; Veloza and Urbina-Cardona 2025). Understanding the response of these patterns in both groups is particularly relevant in tropical montane systems, where environmental gradients and land-use change interact to shape biodiversity.

In heterogeneous landscapes, agroecosystems may play a significant role in conserving biodiversity. Some agricultural systems, such as shaded coffee crops, have been proposed as biodiversity-friendly land uses capable of maintaining a subset of native species while supporting economic production (Perfecto and Vandermeer 2008; Céspedes and Bayly 2019; González et al. 2020; Ríos-Orjuela et al. 2024). These systems can function as intermediate habitats or dispersal matrices between forest remnants. However, despite increasing evidence of the effects of land-use change, there is still limited understanding of how multiple dimensions of diversity and community structure respond simultaneously across heterogeneous landscapes at local scales, particularly in regions with complex mosaics of land use (Brüning et al. 2018; Agudelo-Hz et al. 2019; Roach et al. 2020; Zabala-Forero and Urbina-Cardona 2021).

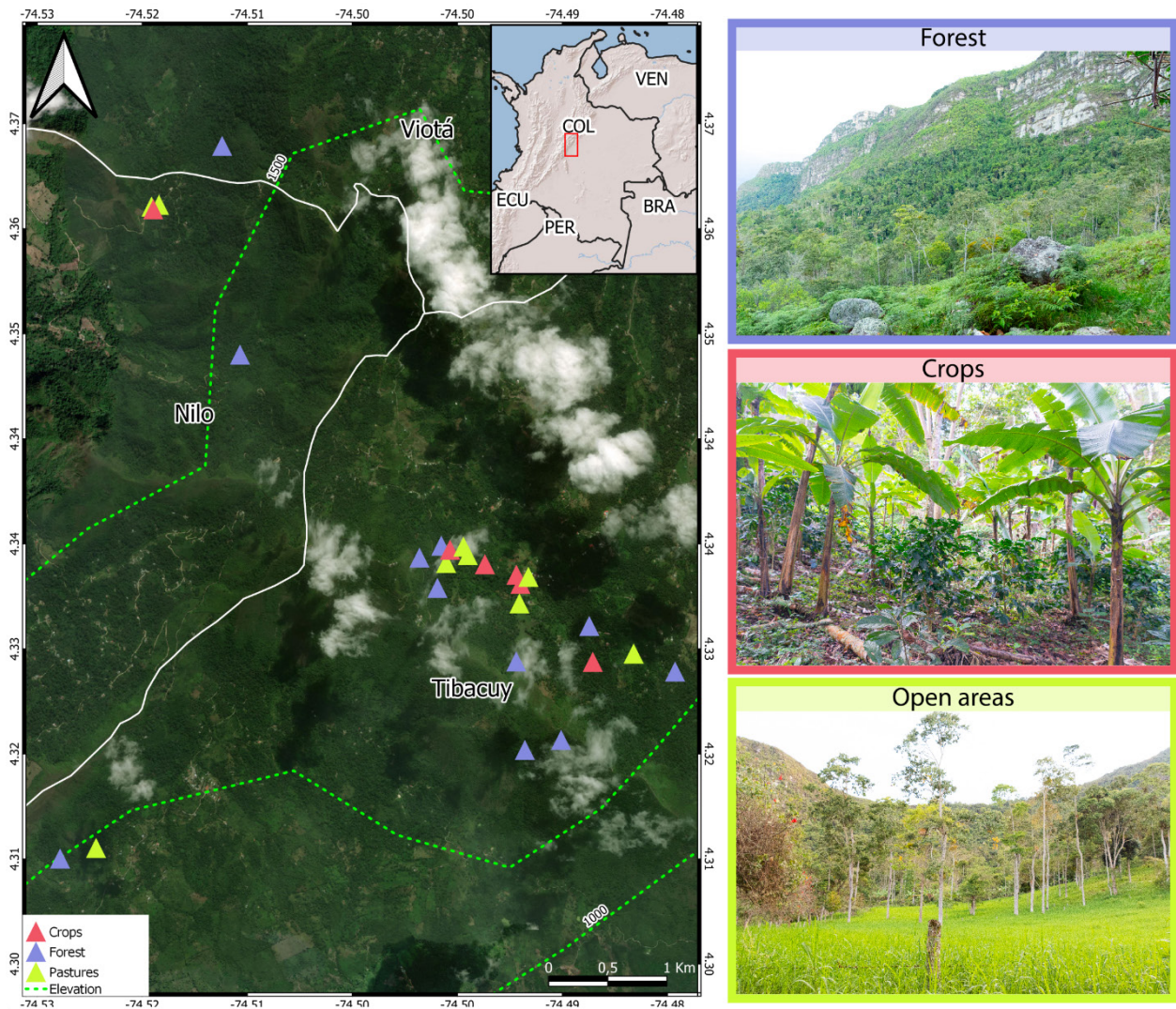
Here, we evaluated how habitat transformation shapes diversity and community structure of amphibians and reptiles across a heterogeneous landscape in the southwestern Cundinamarca region. Specifically, we asked whether changes in habitat structure are associated with shifts in species richness, dominance patterns, and community composition across forest, agricultural, and open habitats. Under the expectation that habitat transformation selectively favors disturbance-tolerant species, we expected that: (1) species richness would decline from forest to more transformed habitats; (2) community structure would shift towards increased dominance and reduced evenness in disturbed habitats; and (3) responses would differ between taxonomic groups, with amphibians more strongly associated with forest environments and reptiles showing broader tolerance to disturbance. These predictions reflect the expectation that habitat transformation selectively favors generalist species while restricting taxa with more specialized ecological requirements.

Rather than directly testing causal mechanisms, our aim is to provide a standardized, coverage-based characterization of diversity and community structure across habitats, generating a robust baseline for understanding how habitat transformation reshapes assemblages. In addition to these analyses, we compiled historical and current occurrence records to generate a comprehensive species inventory for the region, providing a broader biogeographic context for interpreting observed patterns of diversity and community structure.

## Methods

### Study area

We conducted fieldwork in three municipalities of the southwestern Cundinamarca region (Tibacuy, Nilo, and Viotá), located in the Eastern Cordillera of the Colombian Andes, between 645 and 1840 m elevation (Fig. 1). The study area comprises heterogeneous landscapes dominat-



**Figure 1.** Study area and sampling design across a heterogeneous landscape in the southwestern Cundinamarca region in the Eastern Cordillera. Herpetofaunal sampling was conducted across a heterogeneous landscape dominated by forest, crops, and open areas along an elevational gradient, providing the spatial framework for comparing community structure among habitats. The map shows sampling localities in the municipalities of Tibacuy, Nilo, and Viotá (Cundinamarca, Colombia), with points colored by habitat type: forest (blue), crops (red), and open areas (green). Elevational contours (dashed lines) indicate the altitudinal gradient across the study area (645–1840 m). The inset locates the study region within Colombia. Representative photographs illustrate the three habitat types surveyed: forest (top), shaded coffee crops (middle), and open areas (bottom).

ed by secondary Andean forest and agroecosystems with in a mosaic of forest remnants, shaded coffee crops and open areas used for livestock farming (Peña-Torres 2016).

According to Holdridge’s life zone classification (Holdridge 1967) the region includes low-montane rainforest and low-montane dry forest zones characterized by predominantly humid mountain climate in combination with local dry enclaves, typical of the middle lands of the Magdalena basin. Sampling was conducted across three habitat types: secondary forest (11 localities), with scattered trees over five meters high, where advanced plant regeneration processes are evident; shaded coffee crops (6 localities) and open areas (9 localities) used for

permanent livestock. Geographic coordinates, elevation, sampling period, and number of transects per locality are provided in Suppl. material 1.

### Field sampling and sampling effort

Herpetofauna surveys were conducted in two periods: between 2015 and 2019 and between June 2021 and July 2022, covering the rainy and dry seasons in the region. During the sampling phase, we employed time-constrained Visual Encounter Surveys (VES), following standard protocols for amphibian and reptile inventories (Villarreal et al. 2006).

Sampling consisted of active visual and auditory surveys within representative microhabitats including vegetation, leaf litter, streams, and ponds, up to a maximum height of 5 m above ground (Cortés et al. 2008). Auditory detections were estimated by a single observer to ensure consistency and represent approximate counts of calling individuals.

Each sampling unit consisted of a four-hour search conducted by five observers. Surveys were performed during both diurnal (8:00–12:00) and nocturnal (18:00–22:00 h) activity periods. In total, 53 sampling events were conducted across the study area between 2015 and 2022, corresponding to a total sampling effort of 1060 person-hours. Sampling effort was broadly comparable across habitats, although it varied slightly among them, with 500 person-hours in forest habitats, 340 person-hours in crops, and 220 person-hours in open areas. Survey sites were selected within representative areas of each habitat type. Not all localities were visited in every sampling period; however, each sampling event was treated as a sampling unit within habitat categories, as analyses were designed to compare community patterns among habitats rather than to infer independent population-level processes. The distribution of sampling localities, periods of visitation, and number of transects per site are detailed in Suppl. material 1.

## Community analyses

Habitat types were defined based on vegetation structure and land use following a simplified classification scheme. Forest sites correspond to areas with continuous or semi-continuous tree cover and advanced vegetation structure. Coffee crops include shaded or semi-shaded agricultural systems with varying management intensity. Open areas comprise pasture-dominated landscapes, including grasslands with sparse or no tree cover. These categories represent a gradient of habitat transformation from less disturbed (forest) to highly modified environments (open areas). A detailed description of vegetation cover units is provided in Ríos-Orjuela et al. (2024).

To evaluate species richness at standardized sampling efforts and predict species richness beyond our observed sample size, we assessed sampling completeness using coverage curves for the entire community and by habitats, using rarefaction and extrapolation analyses based on Hill numbers with 1000 bootstrap resampling to estimate 95% confidence intervals (CI). We quantified diversity as Hill numbers representing species richness ( $q_0$ ), Shannon diversity ( $q_1$ ), and Simpson diversity ( $q_2$ ) using the *iNEXT* package (Chao et al. 2014; Hsieh et al. 2016).

To evaluate patterns of species dominance and evenness among habitats, we calculated rank–abundance curves using abundance data grouped by habitat type. Amphibians and reptiles were analyzed separately to better illustrate differences in community structure between taxonomic groups. We log-transformed abundance values to improve visualization of rank distributions. In addition, rank–abundance plots were generated using *ggplot2*

(Wickham 2009), and annotated with *ggrepel* to identify dominant species within each habitat.

Finally, to assess potential spatial autocorrelation in species richness among sampling localities, Moran's I statistic was calculated using species richness per locality and corresponding geographic coordinates, defining spatial weights and using a k-nearest neighbor approach with the *spdep* package (Bivand et al. 2026).

Given the descriptive nature of this study, comparisons among habitats are based on coverage-standardized diversity estimates and their associated confidence intervals, rather than formal hypothesis-testing approaches. All ecological analyses were conducted exclusively using field survey data collected between 2015–2019 and 2021–2022. All analyses were performed in R version 4.1.0. (R Core Team 2023).

## Regional diversity compilation

To compile a comprehensive species inventory for the study region, field records were complemented with additional occurrence data obtained from the Biodiversity Information System of Colombia (SiB Colombia) and biological collections. Records were filtered to include preserved specimens and verified institutional observations to minimize potential identification errors. We also reviewed specimen records from the Instituto de Ciencias Naturales (ICN) at Universidad Nacional de Colombia and the amphibian and reptile collections of the Instituto de Investigación de Recursos Biológicos Alexander von Humboldt (IAvH) for the municipalities of Tibacuy, Nilo, and Viotá (Cundinamarca). These supplementary records were used exclusively to compile the regional species list and were not included in ecological analyses. Finally, we followed Frost (2024) and Uetz et al. (2024) as taxonomy classification systems for amphibians and reptiles, respectively.

## Results

### Field surveys

Field surveys conducted between 2015–2019 and 2021–2022 recorded a total of 57 species, representing a substantial component of the known regional diversity (Suppl. material 2). Across all sampling events, a total of 608 individuals were recorded, including 391 amphibians and 217 reptiles. The most abundant amphibian species were *Boana platanera* (62 individuals), *Engystomops pustulosus* (60), and *Pristimantis taeniatus* (55). Among reptiles, the most abundant species were *Gonatodes albogularis* (53 individuals), *Cnemidophorus lemniscatus* (33), and *Hemidactylus frenatus* (17).

Species richness varied among habitat types, with forest supporting the highest richness (45 species), followed by open areas (34) and coffee crops (14). In terms of abundance, forests also contained the greatest number of individuals (225 records), followed by open areas (207), while coffee crops supported considerably fewer individuals (176).

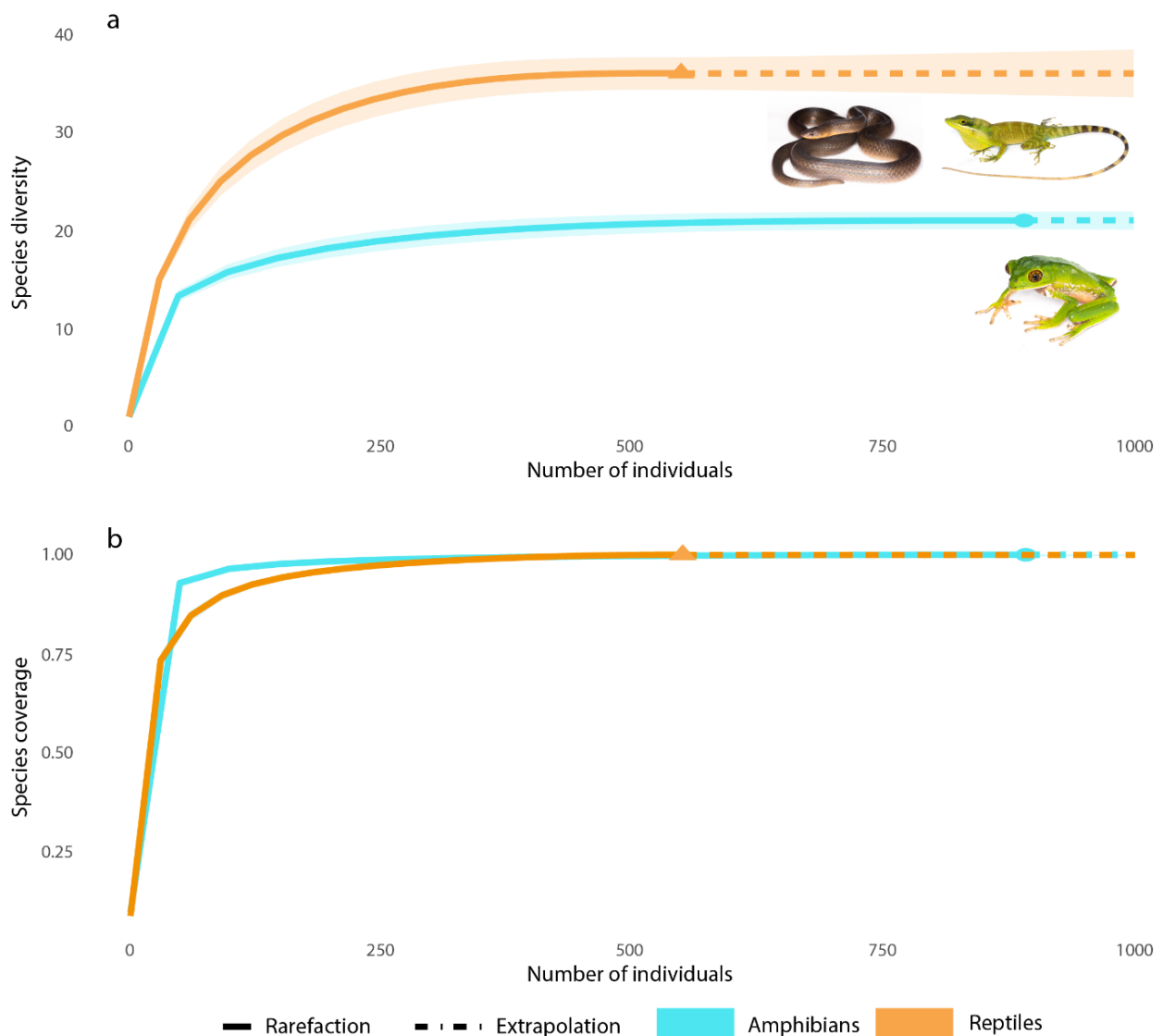
### Sampling completeness

Rarefaction and extrapolation analyses showed that sampling covered a large proportion of the expected diversity of both amphibians and reptiles (Fig. 2a). Sample coverage exceeded 95% for both taxonomic groups (Fig. 2b). Coverage-based rarefaction curves approached asymptotic levels, with amphibians reaching asymptotes at lower sample sizes than reptiles.

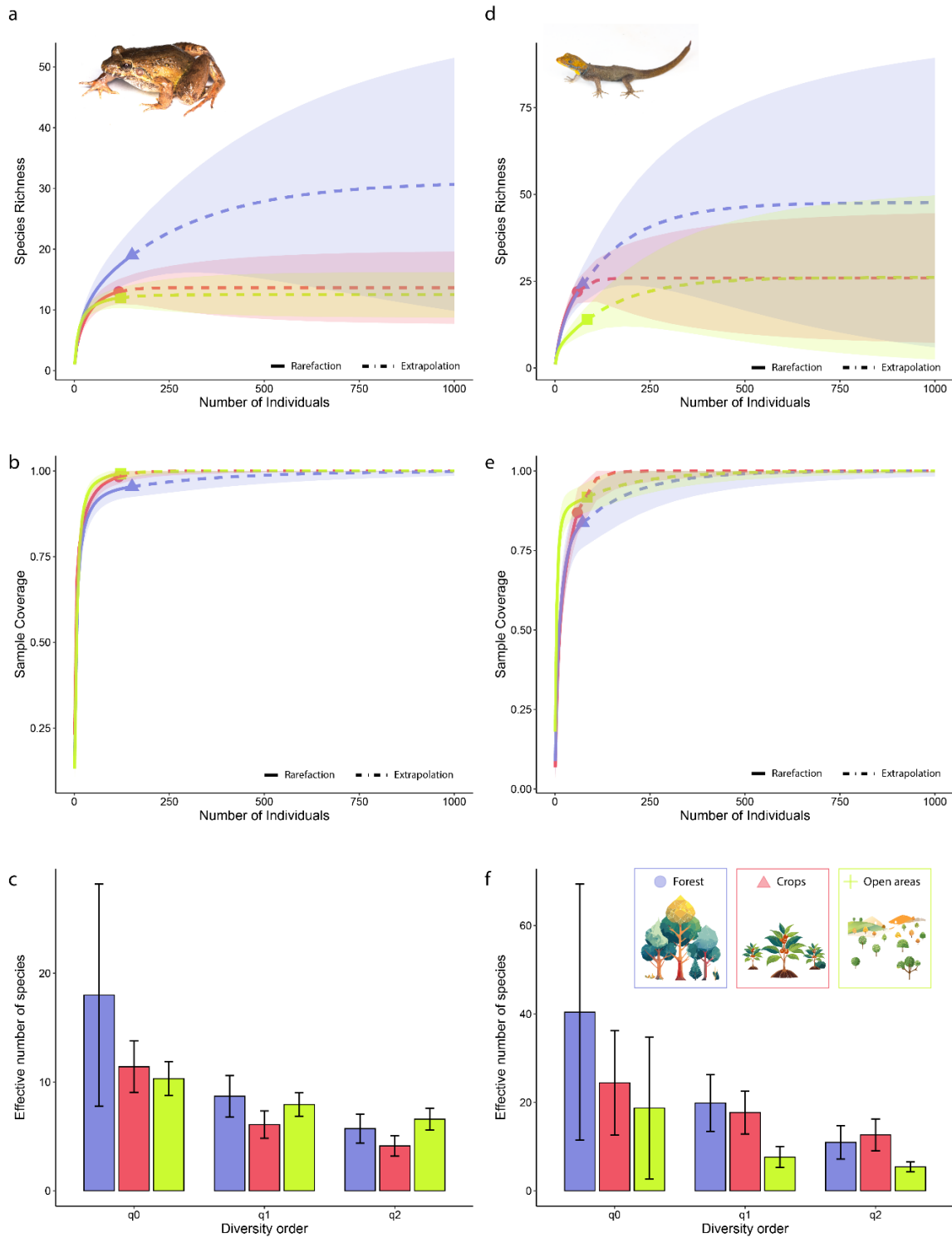
Spatial autocorrelation analysis using Moran’s I revealed no significant spatial structure in species richness across sampling localities (Moran’s I = -0.0147, expected I = -0.040, p = 0.385), indicating that observed richness patterns of the herpetofauna community were not spatially structured.

### Diversity across habitats

Coverage-based rarefaction and extrapolation analyses revealed consistent habitat-driven patterns in diversity for both amphibians and reptiles (Fig. 3). Amphibians exhibited the highest species richness (q0), Shannon diversity (q1) and Simpson diversity (q2) in forest, with reduced diversity in crops and open areas across all Hill numbers. Reptiles showed a similar habitat gradient, with forests supporting the highest diversity; however, overall richness and effective diversity values were lower than those observed for amphibians. In addition, reptile assemblages displayed comparatively stronger reductions in diversity in open areas, particularly in terms of dominance-adjusted metrics.



**Figure 2.** Sampling captured most of the expected diversity, with higher species richness in reptiles than amphibians. Sampling achieved high completeness for both taxonomic groups, and reptiles consistently exhibited higher species richness than amphibians across the full range of observed and extrapolated sample sizes. **a.** Rarefaction (solid lines) and extrapolation (dashed lines) curves show the relationship between species diversity (Hill number q0) and the number of individuals sampled for amphibians (blue) and reptiles (orange). Symbols indicate the observed sample size, and shaded areas represent 95% confidence intervals based on 1,000 bootstrap replicates. **b.** Sample coverage curves show that sampling completeness rapidly approached asymptotic values (>95%) for both groups.



**Figure 3.** Forest habitats support higher diversity, whereas transformed habitats show reduced richness and simplified assemblages in amphibians and reptiles. Species diversity declined from forest to crops and open areas, suggesting that habitat transformation reduces richness and simplifies community structure across the landscape for both groups. **a, d.** Coverage-based rarefaction (solid lines) and extrapolation (dashed lines) curves of species richness (Hill number  $q_0$ ) as a function of the number of individuals sampled for forest (blue), crops (red), and open areas (green) in amphibians and reptiles, respectively. Symbols indicate observed sample sizes, and shaded regions represent 95% confidence intervals based on 1,000 bootstrap replicates. **b, e.** Sample coverage curves show that sampling completeness was similarly high across habitats and taxa, allowing robust comparisons of diversity patterns. **c, f.** Effective diversity (Hill numbers  $q_0$ ,  $q_1$ ,  $q_2$ ) illustrating consistent declines in diversity from forest to more disturbed habitats in both amphibians and reptiles, although amphibians generally exhibit higher richness and diversity values than reptiles. Differences among Hill numbers indicate that habitat transformation affects both species richness and dominance structure, with stronger effects in disturbed habitats. Error bars represent 95% confidence intervals.

When comparing taxonomic groups, both amphibians and reptiles consistently showed higher diversity in forest habitats and reduced diversity under habitat transformation. However, the nature of this response differed between groups: amphibians exhibited a stronger decline in species richness ( $q_0$ ), whereas reptiles showed more pronounced changes in community structure related to species dominance ( $q_2$ ). These patterns suggest that, although habitat transformation affects both groups in a similar way, it impacts different components of diversity in each group.

## Community structure across habitats

Rank–abundance distributions revealed differences in species dominance and community structure among habitats (Fig. 4). Forest assemblages showed longer tails in rank distributions, with a greater number of low-abundance species.

Forest showed higher richness and a more gradual decline in amphibian species abundance, whereas coffee crops and open areas were characterized by fewer species and stronger dominance by a small subset of taxa (Fig. 4b). Dominant amphibian species varied among habitats, including *Engystomops pustulosus*, *Boana platanera*, and *Pristimantis* species. Similarly, the dominant reptile species were *Cnemidophorus lemniscatus*, *Gonatodes albogularis*, and *Hemidactylus frenatus*, whereas forest sites contained a broader distribution of species with lower individual abundances (Fig. 4c).

Overall, these results describe clear differences in community structure across habitat types, with forests showing higher species richness and more even assemblages, and open areas characterized by greater dominance of fewer species.

## Regional herpetofauna checklist

A total of 86 species of herpetofauna have been recorded in the study region based on our field surveys, biological collections, and biodiversity records in databases (Suppl. material 2). Among amphibians, the order Anura is the richest group with 32 of 35 species recorded. The most speciose families were Hylidae (8 species), followed by Strabomantidae and Dendrobatidae (5 species each). Caecilians (Gymnophiona) were represented by two species whereas there are reports of a single species of salamander (Caudata) in the area. Regarding reptiles, 51 species have been recorded primarily represented by the order Squamata (49); the most diverse families were Colubridae (23 species) and Anolidae (6 species). The orders Crocodylia and Testudines were represented by a single species each (*Caiman crocodilus* and *Kinosternon* sp.). Two invasive species were recorded in the region: the North American bullfrog *Aquarana catesbeiana* and the South Asian house gecko *Hemidactylus frenatus*.

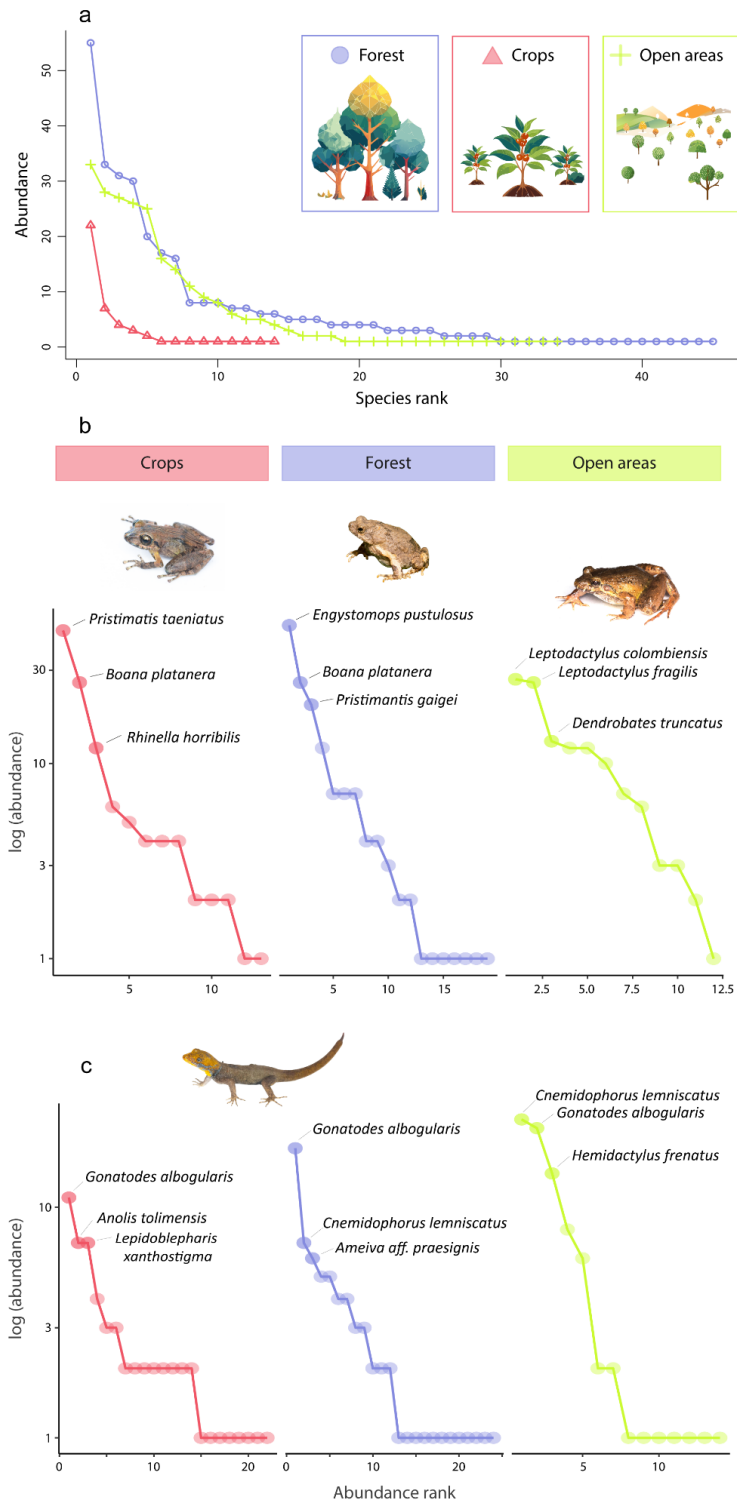
## Discussion

### Drivers of community structure across habitats

Habitat transformation is a major driver of biodiversity patterns in tropical landscapes, affecting both species richness and the internal structure of ecological assemblages (López-Bedoya et al. 2022; Cervantes-López and Morante-Filho 2024). In this study, herpetofaunal assemblages were strongly structured by habitat type, with secondary forests supporting the highest diversity, while agricultural and open areas hosted distinct but simplified assemblages. Despite reduced diversity in modified habitats, a substantial portion of the regional fauna persisted across habitat types, indicating that different components of the landscape contribute unevenly but complementarily to overall diversity.

Although sampling units were spatially connected, the absence of spatial autocorrelation and the consistent differentiation among habitats are consistent with environmental filtering processes associated with habitat transformation. Consistent with this interpretation, habitat transformation is associated with assemblages dominated by generalist species and reduced representation of taxa with more specialized ecological requirements. Similar patterns consistent with environmental filtering have been documented in amphibian and reptile assemblages across transformed tropical landscapes, where changes in vegetation structure, temperature, and moisture favor disturbance-tolerant and generalist species while reducing the representation of forest specialists (Brüning et al. 2018; Zabala-Forero and Urbina-Cardona 2021; López-Bedoya et al. 2022). In Colombia, recent studies have reported that habitat transformation and landscape simplification are associated with changes in the composition of herpetofauna communities through environmental filtering processes that affect species with different ecological requirements, thereby altering the structure of amphibians and reptiles assemblages (Granda-Rodríguez et al. 2025; Palomino-Cuéllar and Urbina-Cardona 2025; Veloza and Urbina-Cardona 2025).

At a regional scale, the southwestern Cundinamarca region harbors a relatively diverse herpetofauna, based on field surveys and compiled records (Lynch 1982; Ramírez 2017; Acosta-Galvis et al. 2020; López-Perilla et al. 2023). Amphibians and reptiles were represented by several families with multiple species, including Colubridae, Anolidae, Hylidae and Strabomantidae, and sampling coverage indicates that the assemblages documented here provide a representative characterization of the local herpetofauna. This diversity reflects the convergence of forest remnants and human-modified habitats across the transition between lowland tropical forests and montane environments, which contributes to shaping regional biodiversity patterns (Brüning et al. 2018; Zabala-Forero and Urbina-Cardona 2021; López-Bedoya et al. 2022).



**Figure 4.** Habitat transformation increases dominance and simplifies community structure, favoring a few abundant species in disturbed habitats. Community structure shifts from more even assemblages in forests to dominance by a few species in transformed habitats, suggesting that habitat modification favors generalist species and reduces community complexity. **a.** Rank–abundance curves for all species pooled by habitat type. Forest (blue) shows a longer tail and more gradual decline in abundance, showing higher evenness and greater representation of rare species. In contrast, crops (red) and open areas (green) exhibit steeper declines, reflecting stronger dominance by a few species. **b.** Rank–abundance curves for amphibians by habitat type (log-transformed abundance). Forest assemblages display greater richness and evenness, whereas crops and open areas are dominated by a reduced number of species. Representative dominant species are labeled for each habitat. **c.** Rank–abundance curves for reptiles by habitat type (log-transformed abundance). Open areas show strong dominance by a few widespread species (e.g., *Cnemidophorus lemniscatus*, *Gonatodes albogularis*, and *Hemidactylus frenatus*), whereas forest assemblages exhibit a more even distribution of abundances across species. Species labels highlight the most abundant taxa in each habitat.

## Diversity patterns across habitat transformation

Clear differences in diversity among habitats reflect variation in structural complexity and microclimatic stability (Cortés-Gómez et al. 2013; Roach et al. 2020; Cervantes-López and Morante-Filho 2024), which promote the persistence of species with specialized ecological requirements and complex life histories (Veloza and Urbina-Cardona 2025). Forest habitats supported the highest diversity across all metrics, whereas crops and open areas exhibited reduced diversity and altered community structure for both amphibians and reptiles.

These patterns align with previous studies indicating that habitat modification reduces species richness and alters community structure in tropical systems (Roach et al. 2020; Galindo-Urbe et al. 2022; Veloza and Urbina-Cardona 2025). Diversity analyses reveal consistent patterns across Hill numbers, suggesting that habitat transformation impacts effective diversity and community structure in amphibians and reptiles (Fig. 3). Forests maintained higher effective diversity and more balanced assemblages, whereas transformed habitats exhibited reduced richness and increased dominance by fewer species. These results suggest that habitat transformation affects the number of species present and the distribution of abundances within assemblages, leading to a simplified community structure across the landscape.

## Contrasting responses of amphibians and reptiles

Differences between amphibians and reptiles may reflect contrasting physiological constraints and ecological strategies. Amphibians were more strongly associated with forest habitats, likely due to their dependence on humid and thermally stable microclimates, which increase the risk of desiccation and reduce reproductive success in disturbed environments (Wanger et al. 2010; Palacios et al. 2013; Méndez-Narváez 2014; Cervantes-López and Morante-Filho 2024).

In contrast, reptiles exhibited broader habitat occupancy, consistent with their higher tolerance to thermal and hydric variation (Méndez-Narváez 2014; Thompson et al. 2015; Cordier et al. 2021; Cervantes-López and Morante-Filho 2024). These differences support the expectation that taxonomic groups respond differently to habitat transformation depending on their physiological and ecological traits.

## Community dominance and structural simplification

Patterns of dominance varied across habitats, with open areas characterized by assemblages dominated by a small number of widespread species (Fig. 4), a pattern frequent-

ly associated with ecological disturbance and habitat simplification (Flynn et al. 2009; Herrera-Montes and Brokaw 2010) and previously reported in the region (Veloza and Urbina-Cardona 2025). In contrast, forest assemblages exhibited a more even distribution of abundances and a greater representation of low-frequency species.

The distribution of species across habitats reflects differences in ecological strategies. Generalist species occurred consistently across all habitats (Suppl. material 2) and tend to dominate local assemblages, indicating high tolerance to environmental variation (Brüning et al. 2018; Zabala-Forero and Urbina-Cardona 2021; Ríos-Orjuela et al. 2024). In contrast, forest habitats harbored species with more restricted ecological requirements, particularly amphibians dependent on stable microclimatic conditions (Galindo-Urbe et al. 2022). These patterns are consistent with a shift toward generalist-dominated assemblages in transformed habitats.

## Role of agroecosystems in heterogeneous landscapes

Agricultural systems may play an important role within heterogeneous landscapes (Galindo-Urbe et al. 2022; Cervantes-López et al. 2025). In the study area, coffee crops hosted several species also present in forest habitats, indicating that these environments can function as secondary habitats or dispersal matrices (Ríos-Orjuela et al. 2024). Agroecosystems therefore may contribute to biodiversity conservation when embedded within heterogeneous landscapes (Roach et al. 2020; Zabala-Forero and Urbina-Cardona 2021; Ríos-Orjuela et al. 2024; Cervantes-López et al. 2025; Pinzón et al. 2025). However, these habitats do not replace forest ecosystems, as they primarily support disturbance-tolerant species and retain only a subset of the regional fauna. Additional dimensions of diversity, such as functional and phylogenetic diversity, should be taken into account to better understand the herpetofaunal assemble dynamics and the role of agroecosystems in conserving community structure.

## Broader implications: land-use change and biotic homogenization

The observed patterns should be interpreted within the broader context of land-use change in the Colombian Andes, where deforestation and agricultural expansion have driven habitat fragmentation (Armenteras et al. 2011; Rodríguez Eraso et al. 2013; Vanegas-Cubillos et al. 2022). These processes tend to favor ecological generalists while reducing the persistence of species with specialized requirements, potentially leading to biotic homogenization and changes in ecosystem functioning (Valencia-Aguilar et al. 2013; Thompson et al. 2015).

In this context, the presence of invasive species such as *Aquarana catesbeiana* and *Hemidactylus frenatus*, which

have been in the region for decades (Rueda-Almonacid 1999; Caicedo-Portilla and Dulcey-Cala 2011; Pérez-Rojas et al. 2026), may further reinforce homogenization processes by favoring disturbance-tolerant assemblages (Rueda-Almonacid 1999; Corporación Autónoma Regional de Cundinamarca CAR 2018; Galindo-Uribe et al. 2022) in a region with one of the highest rates of endemism in amphibians and reptiles (Tobar-Suárez et al. 2022; Díaz-Ricaurte et al. 2025; Vásquez-Restrepo and García-Cobos 2026).

## Limitations and future directions

Our results should be interpreted considering some limitations. Although sampling coverage was high, detectability may vary among taxa and habitats, particularly between amphibians and reptiles. In addition, the elevational gradient encompassed in the study may partially confound habitat effects, as environmental conditions covary with both elevation and land use. Finally, given the descriptive nature of our analyses, the patterns documented here should be interpreted as indicative of underlying ecological processes rather than direct tests of causal mechanisms.

## Regional inventory and taxonomic considerations

In addition to providing ecological insights, this study contributes to improving the knowledge of the regional herpetofauna by presenting the first compiled species list for the southwestern Cundinamarca region in Colombia. Several recorded species represent expected occurrences based on their known distributions in nearby localities but had not previously been documented in detail for the study area. For instance, species such as *Leptodactylus fragilis*, *Rhinella humboldti*, *Cnemidophorus lemniscatus*, and *Bothrops asper* are widely distributed in the Magdalena River valley and surrounding regions, but their occurrence in the surveyed area contributes to refining the regional inventory. These records provide a more robust baseline for future ecological and conservation studies.

There are some records for both amphibians and reptiles worth discussing. Based on currently available morphological information for the species in the region, we were unable to determine one species of *Anolis*. Preliminary observations suggest affinity with *Anolis granuliceps*, although further integrative taxonomic analyses will be required to clarify its identity. The presence of potentially undescribed or poorly characterized taxa highlights the still incomplete knowledge of herpetofauna diversity in the Colombian Andes. Recent studies in the region have revealed several previously unrecognized lineages such as *Anolis tequendama* (Moreno-Arias et al. 2023), *Bolitoglossa muisca* (López-Perilla et al. 2023) and *Hy-*

*loxalus arliensis* (Acosta-Galvis et al. 2020). Likewise, recent systematic studies in *Scinax* suggest that populations currently assigned to this widespread species complexes may correspond to distinct evolutionary lineages (Guarnizo et al. 2015; Araujo-Vieira et al. 2020, 2023).

There are also several relevant historical records in the municipalities studied, especially toward higher altitudes. Species such as *Atelopus subornatus* and *Hyloxalus ruizi* were observed in the area surrounding El Alto de San Miguel by Lynch (1982). Both species are critically endangered (IUCN 2024), and besides the southern records of *A. subornatus* in Tolima (Enciso-Calle et al. 2017), there are no recent records in the area, and we did not register them in this study. Other species currently threatened in this area worth mentioning are *Hyloxalus vergeli*, *Pristimantis uisae* and *P. renjiformis* (Suppl. material 2).

## Conservation implications

Our results underscore the importance of maintaining forest remnants and promoting landscape heterogeneity in the conservation of amphibians and reptiles in the Colombian Andes. Given that habitat transformation is associated with reduced representation of forest-dependent species, continued habitat loss is likely to disproportionately affect taxa with specialized ecological requirements.

Conservation strategies should prioritize the protection of remaining forest patches, the maintenance of ecological connectivity, and the implementation of land-use practices that promote structurally heterogeneous landscapes. Long-term monitoring programs will be essential to evaluate how ongoing environmental changes, including land-use transformation and climate change, affect the distribution and persistence of amphibian and reptile assemblages in Andean ecosystems (López-Bedoya et al. 2022; Cervantes-López and Morante-Filho 2024; Ríos-Orjuela et al. 2024).

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## Supplementary material 1

### Geographic location and total survey events of each location analyzed in the fieldwork

Authors: Nelson Falcón-Espitia, Juan Camilo Ríos-Orjuela, Sebastian Perez-Rojas, Dennys Plazas-Cardona, Alejandra Arias-Escobar

Data type: xlsx

Explanation note: The supplementary material contains the sampling location coordinates and the transect information.

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Link: <https://doi.org/10.3897/herpetozoa.39.e171614.suppl1>

## Supplementary material 2

### Species recorded in the municipalities of the study area (Cundinamarca, Colombia)

Authors: Nelson Falcón-Espitia, Juan Camilo Ríos-Orjuela, Sebastian Perez-Rojas, Dennys Plazas-Cardona, Alejandra Arias-Escobar

Data type: docx

Explanation note: The table displays the species recorded for the study area through fieldwork and database searches.

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Link: <https://doi.org/10.3897/herpetozoa.39.e171614.suppl2>

## Supplementary material 3

### Spanish translation of the manuscript

Authors: Nelson Falcón-Espitia, Juan Camilo Ríos-Orjuela, Sebastian Perez-Rojas, Dennys Plazas-Cardona, Alejandra Arias-Escobar

Data type: docx

Explanation note: The supplementary material is a translation into Spanish of the original manuscript as an effort to make science accessible to all potential readers in the study area.

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Link: <https://doi.org/10.3897/herpetozoa.39.e171614.suppl3>

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### Additional information

#### Conflict of interest

The authors have declared that no competing interests exist.

#### Ethical statement

No ethical statement was reported.

#### Artificial Intelligence (AI) use

The authors accept full responsibility for the content of the manuscript, including the disclosure of any use of AI.

No AI tools were used in the preparation of this manuscript.

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#### Author contributions

Conceptualization: NFE, JCRO. Data curation: AAE, SPR, NFE, JCRO. Formal analysis: NFE, JCRO, SPR. Funding acquisition: AAE, JCRO, NFE, DPC. Investigation: JCRO, SPR, DPC, NFE. Methodology: NFE, JCRO, SPR. Project administration: NFE, JCRO. Visualization: JCRO, NFE. Writing – original draft: JCRO, DPC, NFE, SPR. Writing – review and editing: AAE, NFE, DPC, JCRO, SPR.

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#### Data availability

All of the data that support the findings of this study are available in the main text or Supplementary Information.

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