1 Habitat transformation reshapes diversity and community structure of amphibians

2 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.

Keywords: 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).

One of the main drivers shaping amphibian and reptile assemblages in tropical landscapes is land-use transformation associated with agriculture, livestock, and urban expansion (Ghosh and Basu 2020, Iglesias-Carrasco et al. 2023). These processes lead to habitat loss, degradation, and fragmentation, resulting in heterogeneous landscapes composed of remnant forest patches embedded within human-modified matrices (Thompson et al. 2015, Cervantes-López and Morante-Filho 2024). Such transformations affect not only species richness, but also the composition and internal structure of ecological assemblages, altering

patterns of dominance, evenness, and species interactions (Ernst et al. 2006, Gardner et al.

2007). As a result, disturbed habitats are often characterized by simplified assemblages

dominated by a reduced number of species. However, beyond changes in species richness,

how habitat transformation reshapes the internal structure of assemblages, including

patterns of dominance and evenness, remains less clearly understood.

Differences in species responses to habitat transformation are strongly mediated by

physiological and ecological traits. Amphibians, due to their permeable skin, reproductive

dependence on moisture, 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). In contrast, reptiles tend to exhibit broader tolerance to

thermal and hydric variation, allowing them to persist or even increase in abundance in

disturbed environments (Thompson et al. 2015, Cordier et al. 2021, Veloza and Urbina-

Cardona 2025). These contrasting responses are consistent with environmental filtering

processes, providing a mechanistic framework to understand how habitat transformation

restructures ecological assemblages through shifts in dominance and species composition

(Brüning et al. 2018, Zabala-Forero and Urbina-Cardona 2021, López-Bedoya et al. 2022).

Understanding these patterns 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. Based on environmental filtering theory, 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 dominated by secondary Andean forest and agroecosystems within 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 (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 Supplementary Material 1.

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).

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 Supplementary 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 (q0), Shannon diversity (q1), and Simpson diversity (q2) 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 (Supplementary

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.

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.

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 showed differences in diversity

among habitats (Fig. 3). Forest exhibited the highest estimated diversity across all Hill

numbers with estimated species richness ($q0$) in forests reaching 57.6 species (95% CI:

33.9–81.3), compared to 35.5 species in crops (95% CI: 27.3–43.6) and 23.3 species in

open areas (95% CI: 14.4–32.2). Shannon diversity ($q1$) showed a similar pattern, with

forests exhibiting the highest effective number of species (21.7; 95% CI: 17.4–26.1),

followed by crops (16.6; 95% CI: 14.0–19.2) and open areas (15.2; 95% CI: 12.6–17.8).

Simpson diversity (q2) was similar between forests (11.4; 95% CI: 8.8–13.9) and open

areas (12.2; 95% CI: 10.2–14.2), whereas crops exhibited lower dominance-adjusted

diversity (8.8; 95% CI: 6.9–10.6). Confidence intervals for species richness (q0) did not

overlap between forest and open areas, whereas estimates for other diversity metrics

showed partial overlap among habitats.

When analyzed separately, amphibians and reptiles also showed distinct diversity patterns

across habitats. Amphibian richness was highest in forests and declined in crops and open

areas. Assemblages in open areas showed relatively higher evenness compared to other

habitats. Reptiles followed a similar gradient in species richness, with forests supporting

the highest diversity, while open areas showed lower diversity and stronger dominance

patterns. In crops, reptile assemblages exhibited relatively even distributions of

abundances. Detailed diversity estimates for amphibians and reptiles across habitats are

provided in Supplementary Material 3.

Figure 3. Forest habitats support higher diversity, whereas transformed habitats show reduced richness and simplified assemblages. Species diversity declined from forest to crops and open areas, suggesting that habitat transformation reduces richness and simplifies community structure across the landscape. (a) Coverage-based rarefaction (solid lines) and extrapolation (dashed lines) curves of species richness (Hill number q0) as a function of the number of individuals sampled for forest (blue), crops (red), and open areas (green). Symbols indicate observed sample sizes, and shaded regions represent 95% confidence intervals based on 1,000 bootstrap replicates. (b) Sample coverage curves show that sampling completeness was similarly high across habitats, allowing robust comparisons of diversity patterns. (c) Effective diversity (Hill numbers q0, q1, q2) by habitat type. Forest

exhibits the highest richness and Shannon diversity, whereas crops and open areas show

reduced diversity. Differences in Simpson diversity (q2) suggest comparable dominance

structure between forest and open areas, but reduced diversity in crops. Error bars represent

95% confidence intervals.

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.

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.

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. Open areas were dominated by a few highly abundant species such as *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 (Supplementary

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 suggest that environmental filtering, rather

than spatial dependence, plays a central role in structuring community patterns. Under this

framework, habitat transformation is associated with assemblages dominated by generalist

species and reduced representation of taxa with more specialized ecological requirements.

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).

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.

These patterns are consistent with previous studies showing that habitat modification

reduces species richness and modifies community structure in tropical systems (Roach et al.

2020, Galindo-Uribe et al. 2022, Veloza and Urbina-Cardona 2025). In particular, the non-

overlapping confidence intervals observed between forest and open areas for species

richness highlight a marked difference in diversity between less disturbed and highly

transformed habitats, while partial overlap in other diversity metrics suggests more subtle

changes in dominance and evenness.

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 frequently

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 (Supplementary 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-Uribe 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-Uribe 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 *Hyloxalus 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. renjiforum* (Supplementary 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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