1 **Historical shifts, geographic biases, and biological constraints shape mammal**
2 **taxonomy**
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Short running title:
Temporal trends in mammal alpha taxonomy
Moroti *et al.*

Abstract

Species descriptions in taxonomy have become increasingly comprehensive, yet

disparities persist across taxa and regions. We assess temporal trends in mammal species

descriptions (1990–2022) using four proxies of comprehensiveness—counts of examined

specimens and compared taxa, number of pages (only from the Methods/Results

sections), and number of evidence lines (i.e., analytical tools and techniques). Using

generalised linear models, we assessed how these proxies are explained by factors

associated with species' biology, geography, and taxonomic practice. Most new species

derive from tropical regions, particularly rodents and bats, reflecting global discovery

hotspots. Descriptions have grown more rigorous over time, with expanded specimen

sampling, broader taxonomic comparisons, and integrative methods. However, disparities

emerge along geographic and biological axes: temperate-region descriptions incorporate

more evidence lines, while small-bodied and tropical species (especially bats) remain

understudied due to sampling biases and resource limitations. Body size inversely

correlates with description length, as smaller species often require advanced diagnostics.

Species-rich genera show greater comprehensiveness, likely due to heightened diagnostic

scrutiny, though bats exhibit the opposite trend, possibly due to taxon-specific sampling

challenges. Our findings highlight progress in taxonomic rigor but underscore persistent

gaps tied to geography, body size, and accessibility of analytical tools. Addressing these

disparities requires targeted investments in local capacity, equitable collaboration, and

accessible methodologies to strengthen global taxonomic infrastructure and support

conservation priorities.

Keywords: Taxonomy, Linnean shortfall, Mammalia, species descriptions, taxonomic

practice.

Introduction

Species represent fundamental units in biodiversity science (Braby, Hsu, and Lamas 2024), where their names constitute testable hypotheses subject to revision in light of new data (Bremer et al. 1990; Gippoliti, Farina, and Andreone 2024; Raposo et al. 2017; Streicher et al. 2023). Although taxonomic science is dynamic and an absolute nomenclatural stability is clearly undesirable (Dominguez and Wheeler 1997; Dubois 2007), avoiding unnecessary nomenclatural changes in species identities can increase usability of biodiversity databases and improve scientific communication, data sharing, and conservation planning (Bertrand and Härlin 2006; Hamdan et al. 2024; Mahony et al. 2024; Sullivan et al. 2023). Following the development and popularization of integrative taxonomy, contemporary species descriptions often show increasing robustness—i.e., incorporating more comprehensive diagnostic criteria (Guedes et al. 2024; Poulin and Presswell 2016; Sangster and Luksenburg 2015)—and reduced synonymisation rates (Alroy 2002; Baselga et al. 2008; Guedes et al. 2025; Solow, Mound, and Gaston 1995), contrasting with older descriptions, which generally relied on simpler diagnostics (Guedes et al. 2020; Poulin and Presswell 2016; Sangster and Luksenburg 2015). Understanding how temporal trends in species description comprehensiveness are related to improved sampling and analytical frameworks help to inform best practices that can strengthen taxonomic science.

Comprehensive descriptions can help mitigate common pitfalls in taxonomic practices (Braby et al. 2024; Dayrat 2005; Wüster et al. 2021), such as oversplitting due to insufficient diagnostic criteria and failure to accurately distinguish natural variation within species (Isaac, Mallet, and Mace 2004), or undersplitting resulting from overlooked phenotypic or genetic divergence (Chek, Austin, and Lougheed 2003; Hendrixson and Bond 2005; Mutanen et al. 2016). While the accumulated evidence highlights temporal shifts in taxonomic practices (Guedes et al. 2024; Padial and De la Riva 2020; Sangster and Luksenburg 2015), recent studies also reveal pronounced spatial variation (Freeman and Pennell 2021; Padial and De la Riva 2020; De Queiroz 2007). A well-documented latitudinal taxonomic bias exists, wherein biodiversity-rich but resource-limited regions (e.g., the Tropics) often exhibit lower taxonomic resolution compared to temperate, high-income regions (Diniz Filho et al. 2023; Freeman and

Pennell 2021; Guedes et al. 2025). This spatial variation likely emerges from differences in access to resources and tools, though international collaborations and open-access data initiatives have the potential to mitigate these imbalances (Carneiro et al. 2025; Nakamura et al. 2023). Addressing these gaps is critical to ensuring robust taxonomic frameworks that support downstream research on fields dependent on taxonomic data (Diniz Filho et al. 2023; Lessa et al. 2024).

Because citation-based metrics are poor proxies of the comprehensiveness of taxonomic publications (Pyke 2014; Valdecasas 2011; Valdecasas, Castroviejo, and Marcus 2000), more direct metrics are needed to investigate trends in the robustness of species descriptions and to inform whether many nomenclatural changes are likely to occur in the future across organismal groups. Using metrics like page counts and the number of evidence lines or number of techniques adopted (e.g., morphometrics, cytogenetics, internal anatomy, molecular analysis) in descriptions, studies have generally observed an increase in comprehensiveness of descriptions across taxa (e.g., Benton 2008; Guedes et al. 2024; Pante, Schoelinck, and Puillandre 2015; Poulin and Presswell 2016; Sangster and Luksenburg 2015). Here, we investigate how several proxies of comprehensiveness in species description have changed over time and assess their potential determinants.

We focus our assessment on recent mammals, a group with a well-established alpha taxonomy. Although future additions to mammalian species counts may be small compared to vertebrates like amphibians and reptiles (Moura and Jetz 2021), there are still likely hundreds of species awaiting discovery and description (Parsons et al. 2022). We explore whether temporal variation is evident across four proxies of comprehensiveness in species descriptions, namely: (1) the number of specimens used to describe the new taxa, (2) count of taxa to which the new species is compared, (3) page counts, and (4) number of evidence lines used in the description. We examine whether variability in these proxies is related to species' biology, geography, and taxonomic practice. Additionally, we inspect how accessibility to specific resources and tools is associated with the internationalization of species descriptions. Given that many mammalian taxa still lack comprehensive taxonomic studies (Feijó and Brandão 2022), and that most formal descriptions represent the splitting of one species into many (Burgin et al. 2018; Isaac et al. 2004), it is likely that several taxonomic changes lie ahead. Our

findings can inform the development of protocols and practices that ultimately increase
species description comprehensiveness and improve mammal data usage in biodiversity
research (Mahony et al. 2024).

Material and methods

Data collection

We examined the descriptions of 1,032 mammal species published between 1990 and
2022, covering 33 years of mammal discovery. Focusing on post-1990 descriptions
minimize inconsistencies from historical variations in taxonomic standards, such as the
cladistics and molecular systematics, ensuring comparable scientific and socioeconomic
trends. We only considered "original" descriptions, disregarding redescriptions,
revalidations, and other taxonomic works. Taxonomic nomenclature follows the
December 2022 version of the Mammal Diversity Database (MDD 2022).

For each species in our database, we extracted four proxies for the species
description comprehensiveness: (1) number of specimens - as type-specimens plus all
referred specimens of the new species used to obtain diagnostic traits (type-series size
and all referred specimens were highly correlated, Pearson's $r = 0.60$, $p < 0.001$); (2)
number of taxa compared - the count of close-related species directly compared with the
new species; (3) number of pages - by dividing each page into quadrants and including
only the Methods/Results sections. For studies including multi-species
descriptions/redescriptions, we divided page counts by the number of species
described/redescribed. Our fourth proxy was represented by the (4) number of evidence
lines, scored from only one line of evidence to nine lines of evidence based on the
presence or absence of the analysed character in the formal description: (i) dentition (e.g.
dental formula, tooth shape, etc), (ii) internal soft anatomy (e.g. internal morphology of
the stomach, glans, etc.), (iii) external qualitative morphology (e.g. comparison of head
and tail shapes), (iv) microscopic-based trichology, (v) coloration patterns, (vi)
karyotype, (vii) external morphometry (e.g., head-body length, hind foot length, forearm
length), (viii) osteological morphometry (from teeth, skull, mandible and other skeleton
parts), and (ix) DNA (e.g. allozyme, mtDNA, nucDNA, multiLoci). These proxies

showed low multicollinearity (Variance Inflation Factor, VIF < 1.5; Table S1) and limited

pairwise correlations (Spearman's *r* correlation range = 0.03-0.37; Fig. S1), indicating

they capture relatively distinct aspects of description comprehensiveness.

To represent potential determinants of species description comprehensiveness, we

used the seven variables: (1) year of species description; (2) number of authors; (3)

average number of countries per author, the country counts derived from authors

affiliations were divided by author counts to avoid sampling effects associated with

multiauthored works (see Fig. S2 for pairwise correlation between authors and countries

counts); (4) maximum body mass, obtained from original descriptions and from Moura

et al. (2024), (5) per-genus species richness based on the year of species descriptions (i.e.,

for each species, we tallied the total number of currently accepted species described in

the same genus until their year of description); (6) absolute latitude of type-locality; and

whether description was based on (7) taxonomic revision, which was considered as such

only if the work explicitly mentioned the term "revision" (or similar) in its title, abstract,

keywords, or main text. To allow cross-group comparisons, we also recorded data on

taxonomic ranks.

Statistical analyses

Temporal trends in our proxies of the species description comprehensiveness (number of

lines of evidence, number of pages, number of specimens, and number of taxa compared)

were investigated using Spearman's (r_s) correlation between the year of publication and

annual means for the metrics analysed across species described in each year. We assessed

multicollinearity among continuous predictors using the Variation Inflation Factors (VIF;

Mansfield and Helms 1982), where strong multicollinearity is often attributed to variables

holding VIF > 10 (Kutner et al. 2005). All variables in our analysis had VIF values below

2 (Table S2), so none were excluded. We inspected the multicollinearity of the predictor.

We assessed potential determinants of species description comprehensiveness,

using generalised linear models (GLMs; Zuur et al. 2009). We used a negative binomial

error distribution to analyse the number of specimens and number of taxa compared, both

of which are count data. The number of pages was calculated as a continuous variable, so

we applied a $\log_{10}$-transformation and modelled it using a GLM with a Gaussian error

distribution. Although the number of evidence can be considered a count variable, given

its hump-shaped distribution (Fig. S1), we used both negative binomial and Gaussian

distributions, comparing model fit and selecting the one with the lowest AIC values

(Burnham and Anderson 2002). Each response variable was modelled against year of

species description, number of authors (log_{10}-transformed), average number of countries

per author (log_{10}-transformed), body mass (log_{10}-transformed), per-genus species

richness (log_{10}-transformed), and occurrence of taxonomic revision. Continuous

predictors were z-transformed before modelling, allowing comparisons of their effect

sizes. Because of varying amounts of missing data among response and predictor

variables, GLMs were performed with datasets differing slightly in the number of

modelled species: number of lines of evidence ($n = 820$ species); number of pages ($n =$

807), number of specimens ($n = 805$); and number of taxa compared ($n = 810$). All

analyses were performed in the R software version 4.3.1 (R Core Team 2023).

All models were fitted using the package *MASS* (Venables and Ripley 2002). We

assessed explained variation through Nagelkerke's pseudo-R^2 for negative binomial

models (response variables 'number of taxa compared' and 'number of specimens') and

the coefficient of determination (R^2) for the Gaussian model (response variables 'number

of evidence' and 'number of pages'), both computed with the R package *performance*

(Lüdecke et al. 2021). We assessed the necessity of phylogenetic regression models by

investigating the phylogenetic autocorrelation of model residuals through Moran's *I*

correlograms, computed across 14 distance classes (Revell 2010). Correlograms were

derived from the average results of 100 fully sampled phylogenies for mammals (Upham,

Esselstyn, and Jetz 2019), with trees trimmed to include only modelled species.

Phylogenetic autocorrelation analyses were conducted with the R packages *phylobase*

(Bolker et al. 2020) and *phylosignal* (Keck et al. 2016).

To further investigate if accessibility of costly analytical tools is reflected across

international collaboration, we applied a Kruskal-Wallis analysis to test whether the

average number of countries per authors differed among descriptions using (i) molecular

biology, (ii) taxonomic review, or (iii) other evidence lines. *Post-hoc* comparisons were

conducted using Dunn's test, with pairwise contrasts adjusted for multiple testing using

the Bonferroni correction method (Bonferroni 1936). All analyses were conducted using

the *rstatix* R package (Kassambara 2023). See Data Availability section for raw data and

R code.

Results

Most mammal species described between 1990 and 2022 were from Africa, South

America, Southeast Asia, and islands near Australia (Fig. 1). The Afrotropical realm

stands out in terms of primate descriptions, whereas the few descriptions from the

Nearctic were mostly from rodents and Eulipotyphla (e.g., hedgehogs, moles, true

shrews). Of the 1032 species described, 40.8% were rodents (n = 421) and 27.1% were

bats (n = 280)—a trend observed globally as well as across most biogeographic realms.

Descriptive statistics for comprehensiveness metrics across all mammals, rodents, bats,

and when considering mammals excluding bats and rodents, are summarized in Table 1.

Figure 1. Global distribution of mammal species described between 1990 and 2022. Each
point on the map represents a species' type locality, while donut charts indicate the
proportion of species described within each taxonomic order. Points are coloured
according to their respective taxonomic order, with rodents in blue, bats in green, and the
different shades of red representing other mammals. To improve visualization, taxonomic
orders holding less than 5% of species descriptions were grouped under 'Other taxa' label
in the donut charts. The total sample size (N) is lower than the 1,032 species described
during this period due to missing type-locality data for some species.

Table 1. Descriptive statistics for metrics of species descriptions' comprehensiveness.
For each group, we report the median and the range (minimum–maximum) for the number
of evidence lines, number of specimens, number of pages, and number of taxa compared.

	All mammals	Non-bats & non-rodents	Bats	Rodents
N. of evidence lines	5 (1–8)	4 (1–7)	5 (1–7)	5 (1–8)
N. of specimens	8 (1–409)	7 (1–345)	7 (1–327)	9 (1–409)
N. of pages	7.3 (0.23–179)	5.5 (0.23–48.2)	8 (0.8–179)	8.1 (1–130)
N. of taxa compared	5 (1–77)	4 (0–46)	5 (1–45)	5 (1–77)

Temporal trends in mammal description comprehensiveness

The annual mean number of evidence lines used in species descriptions increased when
analyzing all mammals ($rs = 0.67$; $p < 0.001$), non-bats & non-rodent mammals ($rs =$
0.45; $p = 0.03$), bats ($rs = 0.60$; $p = 0.001$) and rodents ($rs = 0.53$; $p = 0.007$; Fig 2a).
There was no change in the number of pages over time (Fig. 2b), but an increase in the
number of specimens when considering all mammals together ($rs = 0.47$; $p < 0.05$, Fig.
2c) and non-bats & non-rodent mammals ($rs = 0.45$; $p = 0.03$). The number of taxa
compared also increased for all mammals combined ($rs = 0.59$; $p = 0.001$, Fig. 2d), non-
bats & non-rodent mammals ($rs = 0.51$; $p = 0.01$) and for bats separately ($rs = 0.59$; $p <$
0.002), but not for rodents ($rs = 0.24$; $p = 0.9$). In addition, the incorporation of molecular
analyses in species descriptions has grown over time, with a noticeable increase in the
proportion of taxa described annually using molecular data (Figs S3-S4).

Figure 2. Variation in proxies of mammal species descriptions comprehensiveness over time (1990–2022). Panels show overall trends for all mammals (gray), non-rodents and non-bats (red), bats (green), and rodents (blue). Proxies include the (a) number of evidence used in species descriptions, (b) number of pages in taxonomic description papers, (c) number of specimens examined in species descriptions, and (d) number of taxa compared to distinguish new species. rs denotes the Spearman's correlation between the year of description and annual means for each variable. We applied Bonferroni corrections to p-values to account for multiple comparisons. Vertical lines represent standard errors, and shaded areas around the regression lines represent 95% confidence intervals.

Internationalization of mammal descriptions

We found a temporal decrease in the average number of countries per author for all taxonomic groups, except bats (Fig. 3). In addition, this 'internationalisation indicator' differed across descriptions using particular analytical tools, with studies using molecular biology and taxonomic review showing lower values of average number of countries per

author. In bats and rodents, only molecular-based descriptions involved fewer countries

relative to other lines of evidence.

The use of molecular data in descriptions varies across mammalian orders, with

Primates (105 species), Didelphimorphia, Carnivora, Macroscelidea, Peramelemorphia,

and Artiodactyla frequently incorporating genetic evidence, while Dasyuromorphia (8

species) lacked such data entirely. International collaborations were common in

molecular-based descriptions, except in Lagomorpha, with three descriptions associated

with a single country (Fig. 4a). Taxonomic revisions contributed substantially to new

species descriptions in Pilosa, Monotremata, and Carnivora (> 50%), but were rare in

species-rich groups like Rodentia and Chiroptera and absent in depauperate orders (e.g.,

Peramelemorphia, Paucituberculata, Microbiotheria, Macroscelidea, Lagomorpha, and

Hyracoidea). While international collaboration in revision-based descriptions was

idiosyncratic, primates and afrosoricids exhibited broader multinational involvement

(Fig. 4b).

Figure 3. Average number of countries per author involved in mammal species descriptions across different lines of evidence and over time. Patterns are shown for (a) all mammals, (b) non-bats and non-rodents, (c) bats, and (d) rodents. 'Other evidences' refers to descriptions that do not include molecular diagnoses and were not derived from taxonomic revisions. Molecular approaches exhibit different levels of internationalization compared to taxonomic reviews and other evidence types across mammals, bats, and rodents. In the violin plot, post-hoc comparisons were performed using estimated marginal means, with pairwise contrasts adjusted for multiple testing using the Bonferroni method. Groups that do not share letters are significantly different from each

293 other.
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Figure 4. Proportion of species descriptions using (a) molecular analysis or (b) taxonomic review for each mammal order from 1990–2022. The bars indicate the proportion of the total number of species described per order over this period. The different shades of gray indicate the proportion of these descriptions involving one, two, and three or more countries.

Determinants of comprehensiveness in mammal descriptions

GLMs explained 6-37% of the variation in taxonomic comprehensiveness metrics (Fig.

5). Across all four proxies tested, the number of taxa compared emerged as the best-

explained one for descriptions in all mammals and bats, whereas the number of specimens

was the most explained metric in rodents and non-bats & non-rodent mammals. We did

not find phylogenetic structure in GLM residuals, as indicated by phylogenetic

correlograms (Fig. S5).

We found the year of description, number of species/genus, and absolute latitude

among the most important predictors of species description comprehensiveness. Recent

descriptions used more evidence, referred specimens, compared taxa, and (to some

extent) more pages across all taxonomic groups, although such effect was less

pronounced in bats overall (Fig. 5). Species-rich genera correlated with more specimens

and compared taxa overall, except for bat specimens. Our results revealed a latitudinal

gradient, with the number of evidence lines decreasing in mammal descriptions from
tropical regions. We also observed fewer specimens in descriptions of non-rodent and
non-bat species towards the Tropics.

Other predictors affected only some taxonomic groups. For instance, we found
fewer evidence lines and pages in descriptions of small-sized mammals that are neither
rodents nor bats. In bats, fewer specimens were used to describe small-sized species,
particularly if such descriptions were produced via taxonomic review. However, for non-
flying mammals, the taxonomic reviews tended to promote descriptions with more
evidence lines and to some extent, more pages.

The two remaining predictors, number of authors and average number of countries
per author (an indicator of internationalisation), showed more contrasting trends across
taxonomic groups and proxies of description comprehensiveness. Non-rodent and non-
bat species had fewer lines of evidence and specimens in multi-authored descriptions. In
contrast, bat descriptions with more authors included more evidence, and international
collaborations had limited influence on description comprehensiveness, as descriptions
of non-rodent mammals with multinational authorship consistently included fewer
specimens.

Figure 5. Influence of putative drivers on proxies of mammal species description
comprehensiveness. Standardized coefficients from the generalized linear models fit to

all species and for bats and rodents separately are shown. Horizontal lines denote the 95%
confidence intervals around each coefficient. Predictors with significant effect sizes (p-
value < 0.05) are those whose horizontal lines do not cross 0. Explained variation is
presented (Model R²) as Nagelkerke's pseudo-R² for negative binomial models (Number
of taxa compared and Number of specimens) and as the coefficient of determination (R²)
for the Gaussian model (Number of evidence and Number of pages).
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Discussion

Most new mammal species described between 1990 and 2022 come from tropical regions,

particularly rodents and bats. This pattern aligns with mammal discovery hotspots (Moura

and Jetz 2021; Parsons et al. 2022) and corresponds to the highest diversity of both orders

in these regions (Abreu et al. 2024; MDD 2025). Although the rate of new mammal

descriptions remains slower than in ectothermic tetrapods (Moura et al. 2018), hundreds

of species await formal descriptions (Moura and Jetz 2021; Parsons et al. 2022), with

68.8% increase in recognized species since the 1980s (~65 new species/year; Burgin et

al. 2025). Many additions result from taxonomic splits, reflecting taxonomic uncertainties

often rooted in species descriptions of limited comprehensiveness, likely due to the

historical unavailability or accessibility of tools that are now commonly used in

taxonomic research (Guedes et al. 2024; Poulin and Presswell 2016; Sangster and

Luksenburg 2015). We find a clear temporal progression toward more rigorous practices

in mammal descriptions, including an increased use of multiple evidence lines, expanded

sampling of referred specimens, and broader taxonomic comparisons (Fig. 2). Our results

reveal substantial heterogeneity in determinants of species description

comprehensiveness, likely associated with taxon-specific sampling challenges.

The increase over time in proxies of mammal species descriptions

comprehensiveness has mirrored trends observed across various taxa (Guedes et al. 2024;

Poulin and Presswell 2016; Sangster and Luksenburg 2015). While this pattern aligns

with the principles of integrative taxonomy (Dayrat 2005; Will, Mishler, and Wheeler

2005), it also shows that many taxonomists were already applying integrative approaches

before the formalization of the concept (Valdecasas, Williams, and Wheeler 2008). The

higher comprehensiveness of recent descriptions can be attributed to technological

advancements and reduced costs, which have democratized access to analytical tools

(Guedes et al. 2024; Nakamura et al. 2023; Trisos, Auerbach, and Katti 2021), like micro-

CT scanning, DNA sequencing, and automated platforms for taxonomic analysis (e.g.,
Blackburn et al. 2024; Grenié et al. 2023; Thomer, Twidale, and Yoder 2018). Taken
together, these technological advances have accelerated species descriptions and
expanded the scope and depth of taxonomic knowledge, contributing to a more
comprehensive understanding of biodiversity. Despite improvements in access to
analytical tools, disparities still persist, as evidenced by a greater number of evidence
lines in descriptions from temperate regions (Fig. 5), where wealthy economies
predominate relative to low latitudes.

While descriptions of new mammal species now involve more authors and
countries overall (Fig. S6), the growth in authorship size has outpaced international
collaboration (Fig. 3). Consequently, we find a lower average number of countries per
author in recent descriptions, which is also noticeable across molecular-based
descriptions. These findings may relate to novel standards in authorship criteria of recent
descriptions (Allen, O'Connell, and Kiermer 2019; Lin 2024). For instance, more
collectors of mammal type-specimens have authored recent descriptions (Fig. S7). This
suggests that recent species descriptions may be based on newly collected specimens
obtained through projects led by these collectors, or that there is an increasing recognition
of the contributions made by fieldwork personnel. In addition, the somewhat limited
internationalisation of molecular-based descriptions corroborates DNA sequencing in
taxonomic research, which is nowadays more affordable by researchers in biodiversity-
rich regions in the Global South. For instance, the Human Genome Project (1990-2003)
required 3 US$ billions, whereas advances like next−generation sequencing reduced costs
to less than $1,000 per genome (Schmidt and Hildebrandt 2017; Wetterstrand 2025). This
shift is evident in the rising prevalence of molecular-based diagnoses in mammal
descriptions (Figs S4-S5).

Among the most consistent findings of this study is the higher comprehensiveness
of descriptions made within speciose genera. On the one hand, this finding may reflect a
sampling effect, as species-rich genera inherently necessitate more extensive within-
group comparisons as additional species accumulate over time (Fig. 2). On the other hand,
alpha taxonomy is inherently comparative, and distinguishing closely related species
becomes particularly challenging in species-rich genera (Gaston, Blackburn, and Loder
1995), where greater phenotypic and genetic variation may require more specimens to

accurately delineate species. Surprisingly, bats exhibited the opposite pattern (fewer
specimens in species-rich genera). It is possible that taxon-specific sampling challenges
explain divergent patterns of specimen acquisition. Our dataset shows that 70% of bat
species in the five richest bat genera - *Myotis*, *Murina*, *Rhinolophus*, *Miniopterus*, and
Hipposideros - are echolocating and have body masses below the median of 10 g. This
pattern is likely attributable to size-dependent sampling bias in mist-netting, where mesh
dimensions favour the capture of larger individuals, and the echolocating frequency of
each species may influence the effectiveness of different mist-net types (Ferreira et al.
2021; Francis 1989). Consequently, the species-rich genera are likely under-sampled,
which may explain their limited representation in bat species descriptions. These size-
related constraints in bat sampling align with the higher frequency of taxonomic revisions
in well-sampled (mostly large-sized) species, with less than one-fifth of bats above
median body mass described in taxonomic reviews.

Species descriptions of larger mammals contain fewer evidence lines and are
shorter than those of smaller species, especially in non-flying mammals. While larger
species' morphologies are typically diagnosable without advanced imaging, smaller
species frequently require (Ziegler and Sagorny 2021). Thus, body size shows a negative
relationship with evidence lines, contrasting to its positive associations with species
description probability and data availability (Meiri 2018; Moura and Jetz 2021). The
inverse association between body mass and description length appears to be mediated
through the number of evidence lines. That is, small species demand more pages to
document their additional evidence lines, a pattern supported by the moderate but
significant correlation between page count and evidence lines (Fig. S1). Notably, body
mass influenced specimen availability only in bats, once again reflecting taxon-specific
sampling challenges. Unlike bats, which face size-biased mist-netting limitations, non-
flying mammals (especially rodents) are more efficiently collected through passive
methods like pitfalls and baited live traps (Tobler et al. 2008), with minimal size-related
bias (Bovendorp, MCCleery, and Galetti 2017; Williams and Braun 1983). These
sampling limitations result in fewer specimens of bats relative to non-flying mammals,
as evidenced by GBIF's global holdings of preserved specimens (1.16 million bats versus
3.6 million rodents and 1.89 million other non-flying mammals; GBIF 2025), although

these figures may also reflect the global distribution of known species by order (MDD

2025).

Overall, our study highlights the significant progress in the comprehensiveness of

mammal species descriptions since 1990, driven by technological advancements and the

broader adoption of integrative taxonomic practices. However, persistent disparities

rooted in socioeconomic inequalities, taxon-specific sampling biases, and the broader

taxonomic impediment (i.e., systemic barriers limiting both taxonomic output and end-

user reliability; Ebach, Valdecasas, and Wheeler 2011) continue to shape taxonomic

output. Bats and other small-bodied tropical mammals face compounded challenges, from

methodological constraints to underrepresentation in either molecular diagnoses or

revisionary studies. We reveal that variation in taxonomic practices mirrors well-

documented latitudinal gaps in taxonomic resolution (Freeman and Pennell 2021; Guedes

et al. 2025). Addressing these gaps requires targeted investments in building local

capacity and inclusive collaboration frameworks to ensure more comprehensive species

descriptions (Carneiro et al. 2025; Nakamura et al. 2023; Trisos et al. 2021). In the

absence of such interventions, the interoperability and integration of taxonomic data risk

being delayed (Sandall et al. 2023), thereby undermining subsequent conservation and

research efforts.

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Data Availability Statement

The R-script and raw dataset used in the analyses of this study will be made available at

Zenodo Digital Repository after potential acceptance. Meanwhile, reviewers can access

the data from https://github.com/jhonny-guedes/mammal_desc_trends

Disclosure Statement

The authors report there are no competing interests to declare.

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Figure legends

Figure 1. Global distribution of mammal species described between 1990 and 2022. Each

point on the map represents a species' type locality, while donut charts indicate the

proportion of species described within each taxonomic order. Points are coloured

according to their respective taxonomic order, with rodents in blue, bats in green, and the

different shades of red representing other mammals. To improve visualization, taxonomic

orders holding less than 5% of species descriptions were grouped under 'Other taxa' label

in the donut charts. The total sample size (N) is lower than the 1,032 species described

during this period due to missing type-locality data for some species.

Figure 2. Variation in proxies of mammal species descriptions comprehensiveness over

time (1990–2022). Panels show overall trends for all mammals (gray), non-rodents and

non-bats (red), bats (green), and rodents (blue). Proxies include the (a) number of

evidence used in species descriptions, (b) number of pages in taxonomic description

papers, (c) number of specimens examined in species descriptions, and (d) number of taxa

compared to distinguish new species. rs denotes the Spearman's correlation between the

year of description and annual means for each variable. We applied Bonferroni

corrections to p-values to account for multiple comparisons. Vertical lines represent

standard errors, and shaded areas around the regression lines represent 95% confidence

intervals.

Figure 3. Average number of countries per author involved in mammal species

descriptions across different lines of evidence and over time. Patterns are shown for (a)

all mammals, (b) non-bats and non-rodents, (c) bats, and (d) rodents. 'Other evidences'

refers to descriptions that do not include molecular diagnoses and were not derived from

taxonomic revisions. Molecular approaches exhibit different levels of internationalization

compared to taxonomic reviews and other evidence types across mammals, bats, and

rodents. In the violin plot, post-hoc comparisons were performed using estimated

marginal means, with pairwise contrasts adjusted for multiple testing using the

Bonferroni method. Groups that do not share letters are significantly different from each

other.

Figure 4. Proportion of species descriptions using (a) molecular analysis or (b) taxonomic

review for each mammal order from 1990–2022. The bars indicate the proportion of the

total number of species described per order over this period. The different shades of gray

indicate the proportion of these descriptions involving one, two, and three or more

countries.

Figure 5. Influence of putative drivers on proxies of mammal species description

comprehensiveness. Standardized coefficients from the generalized linear models fit to

all species and for bats and rodents separately are shown. Horizontal lines denote the 95%

confidence intervals around each coefficient. Predictors with significant effect sizes (p-

value < 0.05) are those whose horizontal lines do not cross 0. Explained variation is

presented (Model R^2) as Nagelkerke's pseudo-R^2 for negative binomial models (Number

of taxa compared and Number of specimens) and as the coefficient of determination (R^2)

for the Gaussian model (Number of evidence and Number of pages).