

Title: Amphibian fungal panzootic causes catastrophic and ongoing loss of biodiversity

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15 Abstract:

Anthropogenic trade and development have broken down dispersal barriers, facilitating the spread of diseases that threaten Earth's biodiversity. We present a global, quantitative assessment of the amphibian chytridiomycosis panzootic, one of the most impactful examples of disease spread, and demonstrate its role in the decline of at least 501 amphibian species over the last half century, including 90 presumed extinctions. The effects of chytridiomycosis have been greatest in range-restricted anurans, wet climates, and high elevations in the Americas and Australia.

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Declines peaked in the 1980s and only 12% of declined species show signs of recovery, while 39% are experiencing ongoing decline. There is risk of further chytridiomycosis outbreaks in new areas. The chytridiomycosis panzootic represents the greatest recorded loss of biodiversity attributable to a disease.

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One Sentence Summary:

The amphibian disease chytridiomycosis has driven the greatest loss of biodiversity attributable to a pathogen.

10 Main Text:

Highly virulent wildlife diseases are contributing to the Earth's sixth mass extinction (1). One of these is chytridiomycosis, which has caused mass amphibian die-offs worldwide (2, 3). Chytridiomycosis is caused by two fungal species, *Batrachochytrium dendrobatidis* (discovered in 1998, 4) and *B. salamandrivorans* (discovered in 2013, 5). Both *Batrachochytrium* species likely originated in Asia and their recent spread has been facilitated by humans (5, 6). Twenty years after the discovery of chytridiomycosis, substantial research has yielded insights about its epidemiology (2, 3, 7, 8), yet major knowledge gaps remain. First, the global extent of species declines associated with chytridiomycosis is unknown (see 2, 9 for initial assessments). Second, although some regional declines are well studied, global spatial and temporal patterns of chytridiomycosis impacts remain poorly quantified. Third, ecological and life history traits have been examined only for a portion of declined species (10, 11). Finally, following initial declines, it is unknown what proportion of declined species exhibit recovery, stabilize at lower abundance,



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or continue to decline. Here we present a global epidemiological analysis of the spatial and temporal extent of amphibian biodiversity loss caused by chytridiomycosis.

We conducted a comprehensive examination of evidence from multiple sources, including the IUCN Red List of Threatened Species (*12*), peer-reviewed literature, and consultation with amphibian experts worldwide (data S1). We classified declined species into five decline severity categories corresponding to reductions in abundance. Species declines were attributed to chytridiomycosis based on diagnosis of infection causing mortalities in the wild, or if this was unavailable, evidence consistent with key epidemiological characteristics of this disease. Most evidence is retrospective because many species declined before the discovery of chytridiomycosis (data S1).

We conservatively report chytridiomycosis has contributed to the decline of at least 501 amphibian species (6.5% of described amphibian species; Figs. 1, 2). This represents the greatest documented loss of biodiversity attributable to a pathogen, and places *B. dendrobatidis* among the most destructive invasive species, comparable to rodents (threatening 420 species) and cats (*Felis catus*) (threatening 430 species) (*13*). Losses associated with chytridiomycosis are orders of magnitude greater than for other high-profile wildlife pathogens, such as white-nose syndrome (*Pseudogymnoascus destructans*) in bats (six species) (*14*) or West Nile virus (*Flavivirus sp.*) in birds (23 species) (*15*). Of the 501 declined amphibian species, 90 (18%) are confirmed or presumed extinct in the wild, with a further 124 (25%) experiencing a >90% reduction in abundance (Figs. 1, 2). All except one species' decline (*Salamandra salamandra* affected by *B. salamandrivorans*) were attributed to *B. dendrobatidis*.

Declines were proportional to taxonomic abundance, with anurans having 93% of severe declines (they comprise 89% of all amphibian species). Within anurans, there has been marked



taxonomic clustering of declines, with 45% of severe declines and extinctions occurring in the Neotropical genera *Atelopus*, *Craugastor* and *Telmatobius* (Fig. 2) (*16*). Chytridiomycosis is lethal to caecilians (*17*), but there have been no caecilian declines due to the disease, although data are limited. The capacity for *B. dendrobatidis* to cause major declines is attributable to its maintenance of high pathogenicity (*2*, *18*), broad host range (*8*), high transmission rate within and among host species (*2*, *7*), and persistence in reservoir host species and the environment (*19*). For many species, chytridiomycosis is the principal driver of decline, exemplified by precipitous mass mortalities in undisturbed environments (*2*). In other species, chytridiomycosis acts in concert with habitat loss, altered climatic conditions, and invasive species to exacerbate species declines (*20*).

Most amphibian declines have occurred in the tropics of Australia, Mesoamerica and South America (Fig. 1), supporting the hypothesis that *B. dendrobatidis* spread from Asia into the New World (6). Asia, Africa, Europe and North America have had remarkably low numbers of declines attributable to chytridiomycosis, despite widespread occurrence of *B. dendrobatidis* (8). Relative lack of documented declines could reflect less knowledge of amphibian populations in Asia and Africa (3, 21), early introduction and potential coevolution of amphibians and *B. dendrobatidis* in parts of Africa and the Americas (e.g. (22)), the comparatively recent emergence of *B. dendrobatidis* in Western and Northeast Africa (6), or unsuitable conditions for chytridiomycosis. It remains unknown whether chytridiomycosis contributed to widespread amphibian declines reported in North America and Europe in the 1950-1960s (3, 21, 22) or current enigmatic salamander declines in eastern North America. While the number of new declines has now eased (Fig. 3), additional declines could occur if *B. dendrobatidis* or *B. salamandrivorans* are introduced into new areas, highly virulent lineages are introduced into

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areas that currently have less virulent lineages (6), and/or environmental changes alter previously stable pathogen-host dynamics (3).

Chytridiomycosis-associated declines peaked globally in the 1980s, between one and two decades prior to the discovery of the disease (Fig. 3, and table S1), and coincident with anecdotal recognition of amphibian declines in the 1990s (23). A second, smaller peak occurred in the early 2000s, associated with an increase in declines in western South America (Fig. 3, and fig. S1). Regionally, temporal patterns of decline are variable (fig. S1). For example, in some areas of South America and Australia, declines commenced in the late 1970s (2, 24), while in other areas, declines started in the 2000s (25). *Batrachochytrium dendrobatidis* is associated with ongoing declines in 197 assessed species. Ongoing declines after a transition to enzootic disease dynamics (19) might be driven by a lack of effective host defences, maintenance of high pathogenicity (18), and presence of *B. dendrobatidis* in amphibian and non-amphibian reservoirs (7, 19).

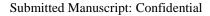
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We examined host life-history traits and environmental conditions to understand why some species declined more severely than others, using multinomial logistic regression and accounting for the degree of evidence that chytridiomycosis was implicated in each species' decline (fig. S2 and table S2). Decline severity was greatest for larger-bodied species, those occurring in consistently wet regions, and those strongly associated with perennial aquatic habitats. These patterns are likely due to favourable environmental conditions for *B*. *dendrobatidis* in wet regions (7), because the fungus dies when desiccated, as well as the general pattern of increased time to maturity in large bodied amphibians resulting in less reproductive potential to offset mortality due to chytridiomycosis (26). Declines were less severe for species with large geographic and elevational ranges (Fig. 4), potentially due to the greater chance of



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their range encompassing environmental conditions unfavourable for *B. dendrobatidis* (*3*), and/or information bias, as population extinctions can be assessed with more certainty in restricted-range species. Our results are consistent with previous studies showing the risk of chytridiomycosis is associated with host aquatic habitat use, large body size and narrow elevational range (*10*, *11*).

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Encouragingly, of the 292 surviving species for which population trends are known, 60 (20%) have shown initial signs of recovery. However, recoveries generally represent small increases in abundance of individual populations, not complete recovery at the species level. Logistic regression showed the probability of recovery was lower for species that experienced more recent or more severe declines, for large-bodied or nocturnal species, and for species occurring at higher elevations (fig. S2 and table S3). When holding those predictors of recovery at their mean value, the chances of a species recovering from a severe (> 90%) decline were less than one in ten. Low probability of recovery for high elevation species might be related to suitable climatic conditions for fungal persistence, as well as limited connectivity to source populations, and/or longer host generation time (26). Some recoveries may be underpinned by selection for increased host resistance (18), while management of concurrent threats may have facilitated other recoveries (a promising avenue for conservation interventions) (27). Unfortunately, the remaining 232 species have shown no signs of recovery.

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The unprecedented lethality of a single disease affecting an entire vertebrate class highlights the threat from the spread of novel pathogens in a globalized world. Global trade has recreated a functional Pangaea for infectious diseases in wildlife, with far reaching impacts on biodiversity (this study), livestock (28), and human health (29). Effective biosecurity and an immediate reduction in wildlife trade are urgently needed to reduce the risk of novel pathogen



spread. As mitigation of chytridiomycosis in nature remains unproven (*30*), new research and intensive monitoring utilizing emerging technologies is needed to identify mechanisms of species recovery and develop novel mitigation actions for declining species.



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Supplementary Materials:

Materials and Methods

Figure S1. Timing of declines by geographic area.
 Figure S2. Multiplicative effect sizes of predictors of decline severity and recovery.
 Table S1. Summary of modelling results for the timing of species declines.
 Table S2. Summary of modelling results for predictors of decline severity.



Table S3. Summary of modelling results for predictors of recovery.

Table S4. Complete legend of taxonomic groups as indicated in Figure 2 (main text).

Data S1 (Excel format)

References (31-47)



Fig. 1. Global distribution of chytridiomycosis-associated amphibian species declines. Bar
plots indicate the number of declined species, grouped by continental area and classified by
decline severity. Brazilian species are plotted separately from all other South American species
(South America W); Mesoamerica includes Central America, Mexico and the Caribbean Islands;
Oceania includes Australia and New Zealand. No declines have been reported in Asia. [Photo
credits (left to right): *Anaxyrus boreas*, Chris Brown; *Telmatobius sanborni*, Ignacio De la Riva; *Atelopus varius*, Brian Gratwicke; *Cycloramphus boraceiensis*, L. F. Toledo; *Cardioglossa melanogaster*, Mareike Hirschfeld; *Salamandra salamandra*, Didier Descouens; *Pseudophryne corroboree*, Corey Doughty].

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Fig. 2. Taxonomic distribution of chytridiomycosis-associated amphibian declines. Each bar represents one species, color denotes the severity of its decline. Concentric circles indicate, from inner to outer, order (Caudata or Anura), family and genus. Full names given only for families and genera including >5 and >2 species respectively; details for all taxa in table S4. Within each taxonomic level, sublevels are ordered alphabetically. Protruding bars indicate species for which there is evidence of recovery. [Photo credits (left to right): *Telmatobius bolivianus*, Ignacio De la Riva; *Atelopus zeteki* and *Craugastor crassidigitus*, Brian Gratwicke].

Fig. 3. Timing of chytridiomycosis-associated amphibian declines. In (A), bars indicate the cumulative number of declines in a given year, stacked by decline severity. For species in which the exact year of decline is uncertain, the figure shows the middle year of the interval of uncertainty, as stated by experts or inferred from available data. In (B), curves indicate the



cumulative number of declines in each decline severity category in time. Arrows mark the discovery of chytridiomycosis in 1998.

Fig. 4. Severity of chytridiomycosis-associated amphibian declines in relation to the

5 geographic (A) and elevational (B) range of species. In (A) each dot indicates a species, located randomly along the perimeter of a circle with radius equal to the log¹⁰ of the species' geographic range in km². In (B), horizontal bars, boxes and vertical bars indicate, respectively, mean, first and second quartiles, and 95% quantiles of elevation ranges within each category of decline severity.





Supplementary Materials for

Amphibian fungal panzootic causes catastrophic and ongoing loss of biodiversity

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This PDF file includes:

Materials and Methods

Figure S1. Timing of species declines by geographic area.

Figure S2. Multiplicative effect sizes for predictors of severity and recovery.

5 Table S1. Summary of modelling results for the timing of species declines.

Table S2. Summary of modelling results for predictors of decline severity.

Table S3. Summary of modelling results for predictors of recovery from declines.

Table S4. Complete legend of taxonomic groups as indicated in Figure 2 (main text).

10 Other Supplementary Materials for this manuscript include the following:

Data S1 (Excel format)



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Materials and Methods

Compilation of declined-species dataset

Chytridiomycosis and amphibian declines

Chytridiomycosis is a lethal disease of amphibians that was discovered in 1998 (1). In amphibians, chytridiomycosis can develop from infection with one of two fungal species, *Batrachochytrium dendrobatidis* (described in 1999, 2) or *B. salamandrivorans* (described in 2013, 3). We identified chytridiomycosis-associated declines caused by infection with either *B. dendrobatidis* or *B. salamandrivorans*. We found evidence for the decline of only one amphibian species (*Salamandra salamandra*) associated with infection with *B. salamandrivorans*, and subsequently for convenience in the following Supplementary Materials, we focus on chytridiomycosis-associated declines resulting from infection with *B. dendrobatidis*.

Two previous efforts have attempted to quantify the number of amphibian species that have experienced declines associated with chytridiomycosis. First, the International Union for the Conservation of Nature (IUCN) Global Amphibian Assessment (GAA), conducted from 2001 to 2004, concluded that 202 amphibian species had experienced "enigmatic" declines, but did not directly attribute these declines to chytridiomycosis (*4*). However, the GAA was conducted when knowledge of chytridiomycosis and its impacts on amphibians was rudimentary. Building on the GAA, Skerratt et al. (*5*) used multiple lines of evidence to argue that *B*. *dendrobatidis* was the agent driving "enigmatic" declines identified in the GAA.

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Since the 2001–2004 GAA, *B. dendrobatidis* has continued to spread into new regions (6) and understanding of chytridiomycosis and its impacts on amphibian hosts has greatly increased. For example, a search for "*Batrachochytrium dendrobatidis*" yields 40 results on the



Web of Science from 2004 and earlier, but 1,258 results for 2005–2017. However, despite these advances, the scale and number of global amphibian declines associated with chytridiomycosis has yet to be empirically quantified, or examined in light of major developments in our understanding of this disease.

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Identifying declined species

We developed an expert-curated list of species that have experienced declines associated with chytridiomycosis (a method analogous with the IUCN approach to evaluating conservation status). We used a two-step process to identify declined species. First, at a continental scale, we searched for evidence of amphibian declines associated with chytridiomycosis. This involved searching the scientific literature and discussions with amphibian experts. Second, if chytridiomycosis-associated declines were identified for a particular continent, we then engaged regional amphibian experts, generally at the country level, to thoroughly compile information on all chytridiomycosis-associated amphibian declines, using the framework and methods described below. Regional experts were chosen based on their professional experience in amphibian ecology in the region, and knowledge of chytridiomycosis. In the case of Asia, there was no evidence for chytridiomycosis-associated declines, and as such, further assessments of amphibians from that continent were not conducted. We acknowledge that this approach could fail to identify some very recent or only locally known declines. However, such cases are unlikely to substantially affect our analyses as existing knowledge would likely be insufficient to determine the role of chytridiomycosis in the decline.

Our focus was on identifying species that had experienced declines due to chytridiomycosis (as specified below), rather than on identifying species in which infection with



either *B. dendrobatidis* or *B. salamandrivorans* has been documented. This is an important distinction, as some amphibian species can be infected with either pathogen, but not experience morbidity or mortality (asymptomatic) or population declines (7, 8). The global database *B. dendrobatidis*-maps [http://www.bd-maps.net] provides information on species in which infections have been documented.

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For the second step of our assessment conducted in 2017, we assembled a team of 41 researchers with expertise in both amphibian ecology and chytridiomycosis to undertake country-level assessments for 24 countries where there was evidence that chytridiomycosis has contributed to amphibian declines. Following the methodology of Scheele et al. (9), for each species, experts collated all available peer-reviewed articles, government technical reports, theses, conference proceedings, books, information from IUCN assessments, and unpublished data on species' status and trends recorded by amphibian ecologists. The scientific literature was searched through Google Scholar and ISI Web of Knowledge. Bibliographies of relevant papers were checked to identify further literature. Relevant information and sources underpinning the inclusion of each declined species in the dataset is provided in data S1. Prior to commencing their assessments, each expert thoroughly discussed the assessment methodology with the lead author (B.C.S.) to ensure consistent interpretation and implementation of the assessment framework.

As initial outbreaks of chytridiomycosis occurred in many regions before its discovery, and many impacted amphibian species are difficult to observe and/or occur in remote habitats, quantifying the role of chytridiomycosis in declines is, for many species, challenging and retrospective. As such, we developed an epidemiological framework (see Metadata tab in data S1) to integrate all available evidence (including laboratory, experimental and field data) to



evaluate the severity, distribution, and timing of amphibian declines caused by chytridiomycosis. This approach of incorporating multiple lines of evidence is commonly used in health disciplines (*10*), has previously been applied to a continent-wide assessment of chytridiomycosis-associated declines (*9*), and uses published criteria (*5*) for determining disease causation in population declines.

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We identified species that had undergone declines in abundance caused by chytridiomycosis. We define decline severity as the total estimated reduction (%) in abundance across a species' range, associated with chytridiomycosis. Most of the assessed species have undergone declines in both abundance and range. However, we elected to focus on changes in abundance because, in the context of chytridiomycosis, declines in abundance and range are often conflated due to environmental context influencing disease impacts, with range reductions cumulating from population extinctions driven by reductions in abundance.

Our quantification of *B. dendrobatidis*-associated declines is likely conservative for three main reasons. First, *B. dendrobatidis* has likely caused the decline of many undescribed species, a phenomenon that might be particularly relevant in the Neotropics where there are many undescribed species (*11*). Second, species may have declined due to chytridiomycosis, but there is currently no evidence linking their decline to the disease. In such cases, these species have not been included in our assessment. Third, the true extent of a species' decline associated with *B. dendrobatidis* may be unrecognized. As such, species may be categorized in a less severe decline category because, while there is evidence for the role of *B. dendrobatidis* in the species' decline in parts of its range, declines in other parts of its range have not (yet) been attributed to *B. dendrobatidis*. In addition, declines could be under-documented in species with large ranges, due to logistical challenges associated with monitoring and surveying widely distributed species.

Declined species were grouped into five broad classes of severity: (1) minor decline
<20% in abundance; (2) severe decline >20% but <90%; (3) extreme decline >90%; (4)

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presumed extinct in the wild (no known extant populations, and no individuals detected at known historical locations, but some reasonable doubt that the last individual has died); and (5) confirmed extinct in the wild (as per IUCN listing). Note that the class "Extinct" includes both "Extinct" and "Extinct in the Wild" sensu IUCN. For each declined species, we also assessed whether there was evidence for ongoing population declines due to chytridiomycosis (i.e. populations experiencing continued declines following the initial emergence of chytridiomycosis) (yes, no, unknown, not applicable [for presumed or confirmed extinct species]), and whether partial recovery of declined species had been observed (yes, no, unknown, not applicable [for presumed or confirmed extinct species]). Species were classified as experiencing ongoing decline if chytridiomycosis was causing continuing reductions in abundance after initial declines associated with chytridiomycosis outbreaks. Species were classified as experiencing partial recovery if increases in abundance were reported after initial declines caused by chytridiomycosis outbreaks. We report the year recovery commenced for populations of species subject to regular surveys. For species where signs of recovery have been reported, but the absence of regular surveys prevent identification of when recoveries commenced, we provide the first year that recoveries were reported. No date is provided when precise information is unavailable. We note that a high degree of caution is needed in interpreting information on species recovery. In many cases, declined species have only shown signs of recovery in some populations, and in general still have greatly reduced abundance compared with historical levels. Detailed examples of how declines were categorized for eight species with varying decline severities are provided in the 'Worked examples' worksheet in data S1.

Timing of species declines

Species were assigned either a year of decline commencement (n = 104), or if this was not known, an uncertainty interval during which the decline was thought to have occurred (n =345). If known, experts provided the year that decline commenced. If the year was unknown, experts estimated a range of years during which the decline could have occurred, or provided a year which could be indirectly related to the timing of decline (e.g. the last year the species was observed as abundant, or the first year that observations were made of populations post-decline (12). In such cases, we represented uncertainty by defining an interval of years. Intervals were either provided directly by experts, or where experts provided only a start- or end-year for decline, we conservatively assumed a ten-year interval, respectively, before or after the stated year. For some Mesoamerican species (particularly in Panama), rather than an interval, the contributors provided temporal information about the first detection of *B. dendrobatidis* in different areas of the region, and based on that information, we inferred the period during which decline might have occurred as a five-year interval on both sides of the first known detection of B. dendrobatidis within the range of the species, based on the IUCN distribution maps (13). In total, information about the timing of declines was available for 449 species out of 501 in our dataset. Species for which no data about the timing of declines were available (n = 52) were excluded from later statistical models (Timing of declines and Predictors of declines and recoveries below). In those models, each species was assigned a single year of decline, to avoid overrepresentation of species with longer uncertainty intervals. For species with a two- or threeyear interval, we used the first and middle year respectively. Where the uncertainty interval of the timing of decline was longer than three years, we drew a random year within that interval to avoid systematic bias towards the beginning, mid- or endpoints of the interval.

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Evidence for the role of chytridiomycosis in species declines

For each declined species, assessors evaluated the following: 1) Was *B. dendrobatidis* diagnosed using histopathology and/or PCR to confirm infection during mass die-offs or sudden declines? 2) Did declines coincide with the documented emergence and spread of *B. dendrobatidis* in the region? 3) Is the declined species highly susceptible to *B. dendrobatidis* in laboratory experiments (i.e. >75% individuals die post exposure in the laboratory) or demographic field studies? 4) Did sympatric species decline simultaneously due to *B. dendrobatidis*? For each question, species were assigned 'yes', 'no', or 'no data'. The strength of evidence linking *B. dendrobatidis* to each species decline was then scored from one to four, with four being the strongest evidence of *B. dendrobatidis*-associated declines. One = expert opinion of the assessor only. Two = single line of correlative evidence. Three = multiples lines of correlative evidence. Four = robust before-after decline sampling demonstrating declines were caused by *B. dendrobatidis*.

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Statistical analyses

We used generalized linear models to assess the temporal trend in the overall number of declines, in the severity of those declines, and in the probability of a species' recovery. We also sought to identify attributes of species distributions (e.g. range size, climatic region, elevation) and life-history traits that were associated with decline severity and recovery. We selected these variables based on previous work on extinction risk in vertebrates (*14*), life-history traits previously linked to chytridiomycosis-associated declines (*15-17*), as well as known bioclimatic preference of *B. dendrobatidis* (*7*, *18*, *19*). All models described below were fit in JAGS (*20*),

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using uninformative priors for all parameters, and run for 150,000 iterations on three Markov chains, with a burn-in of 75,000 and a thinning rate of 10. Convergence was assessed by visual inspection of the chain histories and using the *R*-hat statistic.

5 Timing of declines

First, we modelled the temporal trend in the overall number of declines. As outlined above, we associated each species with a year of decline (as explained in section 'Timing of species declines' above). For each year between 1970 (the earliest year in our dataset) and 2016, we counted the number of species that had that year as the stated or inferred year of decline and used this as a Poisson-distributed response variable, with year of decline as a predictor.

We compared three alternative functions: null (constant number, i.e. no temporal trend), linear (constant increasing or decreasing trend in the number of declines) and quadratic (to reflect an initial increase, followed by a decrease in the number of declines). Given the simple structure of the models, we compared models on the basis of the coefficient estimates and discarded terms for which the 95% credible interval of the posterior distribution of the regression coefficient encompassed zero (analogous to failure to reject the null hypothesis with α =0.05 if working with a null hypothesis significance test). Results are summarized in table S1.

We also modelled the severity of individual declines as a function of time. Because severity was a categorical variable, ordered from 1 to 5, we used a multinomial logit link regression with year of decline, defined as above, as a predictor (moderator). The output of the model can be interpreted as the probability that a species falls in a given severity class. Consequently, a positive coefficient for time (year) would indicate that species with later declines were more likely to suffer more severe impacts. To account for the varying degree of

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evidence supporting the decline of each species, we took an approach analogous to a metaanalysis (21). We defined the prior variance of the linear estimator for the multinomial probability of a species falling in a higher severity class as a function of the corresponding amount of evidence (1, 10, 100 and 1000 for the four levels of evidence from highest to lowest see *Evidence for the role of chytridiomycosis in species declines*). We added a random effect to account for taxonomic autocorrelation (using the genus of each species as a blocking factor, nested within its family).

We also modelled species recovery (yes/no) using logistic regression and year of decline as a predictor, following the same procedure as above. Here, a negative regression coefficient would suggest species with earlier declines were more likely to have recovered (as expected if there was a time lag between decline and recovery). We excluded species from this analysis if they had gone extinct or if there was no data on population trajectory after the initial decline, leading to a final set of 254 species (including 60 known recoveries).

- Finally, visual analysis of the dataset also led us to hypothesize that declines (particularly less severe ones) may have been under-reported or supported by less evidence before 1975. Therefore, we repeated all analyses described below selecting only those species for which the exact or inferred year of decline was 1975 or later. All results were confirmed with and without pre-1975 data.
- 20 Predictors of declines and recoveries

We sought to identify factors that explained the severity of declines and the probability of species recovery. With severity of decline (1 to 5) and recovery (yes/no) as the respective response variables, we used the same model formulations as above, respectively, multinomial



and logistic regression. For severity of impact, we again used the level of evidence to weigh the prior variance of the linear predictor. For both analyses, we again added a random effect at the genus and family level to account for taxonomic autocorrelation.

We began the analysis for severity with a full model that included 10 covariates, selected on the basis of data availability, hypotheses about host and pathogen ecology, and preliminary 5 visual analysis of the data (see next paragraph for detailed descriptions). The 10 covariates were: (1) Geographic range, expressed as the \log_{10} of the extent of the species range in km²; (2) Latitude, calculated as the absolute value for the centroid of the species' distribution; (3) Mean elevation across the species range (included because it is associated with a range of factors that may either affect suitability for *B. dendrobatidis*, such as temperature and precipitation, or host characteristics that may affect vulnerability to declines, such as lower connectivity in high elevation species and increased age to maturity (22); (4) Elevational range, expressed as the \log_{10} of the difference between the maximum and minimum elevations across the species' range; (5) Body size, expressed as the \log_{10} of the mean snout-to-vent length for the species; (6) Clutch size, expressed as the \log_{10} of the average number of eggs/offspring for the species; (7) Three variables describing association with aquatic habitat: use of aquatic habitat (1 if adults of the species use aquatic habitats for at least part of the year, 0 for species fully terrestrial as adults), association with permanent water bodies, and direct development (note that because these three variables were highly correlated with each other (Pearson's r>0.5, see below), only one was 20 included in the model at a time); (8) Activity pattern of the species (0 for nocturnal species, 1 for all other species not strictly nocturnal); (9) Maximum temperature of the warmest month, averaged over the species range from BIOCLIM05 variable (23); and (10) Precipitation of the driest quarter of the year, averaged over the species range from the BIOCLIM17 variable (23).

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Another 20 additional variables potentially associated with chytridiomycosis were discarded after preliminary modelling and visual analysis (clear absence of visual patterns when plotted, large credible intervals for regression coefficients centered close to zero), leaving the 10 listed above as the set we analysed in detail. These discarded variables included environmental specialization, expressed as the number of environments (arid, temperate, sub-tropical, tropical, sub-alpine, alpine) where the species is known to occur, habitat topography, additional reproductive modes such as use of bromeliads, lakes, wetlands or terrestrial clutches, all BIOCLIM variables other than BIO05 and BIO17, including maximum and minimum annual temperature and precipitation as well as seasonality patterns. The final set of 10 variables covered a wide range of ecological hypotheses about chytridiomycosis-related declines (see below).

Species distribution maps were obtained from the IUCN Red List of Threatened Species (24), from which the geographical range was calculated in ArcGIS 10.1. Minimum and maximum elevations used to acquire elevational ranges (with median elevation calculated as the midpoint) were obtained from species-specific online IUCN Red List accounts. Body size and clutch size were extracted from the AmphiBIO database (25) and were included to reflect life history strategies. Data about association with aquatic habitat were provided by contributors of our dataset and were included to reflect the previously demonstrated link between *B*. *dendrobatidis*-driven declines and aquatic habitats (*15, 26*); activity patterns of the species were compiled from AmphibiaWeb (27) and Stuart et al. (28), and were included due to the thermoregulatory behavior of some species that can inhibit *B. dendrobatidis* growth (29, 30). Finally, averaged temperature and precipitation data spanning each species' range were obtained from the corresponding BIOCLIM variables of the WorldClim 2 dataset (23) at 30 arc second

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(±1x1 km) resolution, calculated across the IUCN Red List distribution maps in ArcGIS 10.1. We chose the BIOCLIM05 and 17 variables to reflect whether, within a species range, part of the year was especially unsuitable for *B. dendrobatidis* persistence and growth, due to a marked dry and/or warm season. We also evaluated interaction terms between environmental (BIOCLIM) and elevation/range covariates. Finally, we added to the initial model a covariate for year of decline where the previous analysis (see above) suggested a significant relationship between time of decline and severity/recovery. For recovery, given the smaller initial dataset, we fitted the initial model with a subset of these variables, following the rule of thumb of at least ten observations in the less represented class (recovery yes/no) for each predictor, and assessing all possible combinations of covariates within that constraint. Because species that have experienced very large declines are unlikely to recover for demographic reasons, we excluded from the analysis all species known or presumed extinct, for which recovery would be impossible by definition, and included an additional predictor to reflect such small-population bottlenecks (0 if the species suffered a decline > 90%, and 1 otherwise). This allowed us to account for demographic barriers to recovery, and identify environmental and life history correlates of recovery among species that may have remained unaffected by such barriers.

Preliminary analysis confirmed that correlation between pairs of predictors did not exceed r = 0.5, except for the three variables describing association with aquatic habitat, of which only one at a time was included in models. In particular, we noted that median elevation and elevational range of species had a Pearson's correlation of r = 0.01, allowing the inclusion of both variables in the models. We centered all continuous variables by subtracting their mean to improve convergence. Rather than using complex imputation techniques, for each combination of covariates we removed species for which values of at least one covariate were missing. We fit

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the model in JAGS using the settings and convergence diagnostics described above. We then progressively simplified the model by removing covariates for which the 95 % credible interval of the posterior distribution of the regression coefficient encompassed zero (analogous to failure to reject the null hypothesis with $\alpha = 0.05$ if working with a null hypothesis significance test). As a covariate was dropped from the model, we added back to the data set any species that had been removed because they missed a value for that covariate, after confirming there was no taxonomic or geographic bias in species that were added at a later stage. When two or more covariates had to be removed, we removed one at a time, beginning from the covariate with the smallest available sample size. Because the three variables describing association with aquatic habitat were highly correlated, only one was included in the model at a time, so we repeated the entire process for each of them. For decline severity and recovery, respectively, the full models contained 102 and 84 species (mostly reflecting limited information about body size and clutch size). As explained above, in the multinomial logistic regression for severity of decline, a positive coefficient for a given covariate would indicate that species with higher values of that covariate (e.g. larger body size) were more likely to be classified in a higher severity class. In the logistic regression for recovery, a positive coefficient would indicate that species with higher values of that covariate were more likely to have recovered from the decline. Results are summarized in tables S2-S3.

We also evaluated the realized effect of each covariate on the response variable, calculating the multiplicative effect size *E* of covariate *i* as the product of the corresponding estimated regression coefficient β by the range of that covariate in the dataset: $E_i = e^{\beta_i (max_i - min_i)}$. Multiplicative effect sizes indicate how much the odds of the response variable (i.e. being classified in a higher severity of decline or having recovered from decline) increase or

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decrease across the observed range of the covariate: values greater and smaller than one indicate positive and negative effects respectively, while a value of one suggests no effect. Figure S2 illustrates the cumulative effect sizes for all predictors for decline severity (fig. S2 A) and recovery (fig. S2 B).



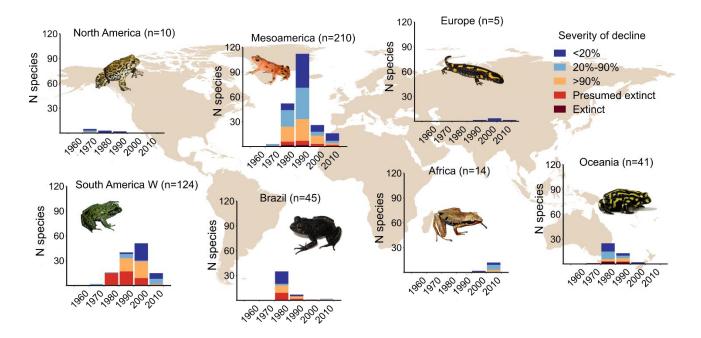


Figure S1. Timing of species declines by geographic area. Bars indicate the number of declines in a given decade, stacked by decline severity. For species in which the exact year of decline is uncertain, the figure uses the middle year of the interval of uncertainty, as stated by experts or inferred from available data. Brazilian species are plotted separately from all other South American species (South America W); Mesoamerica includes Central America, Mexico and the Caribbean Islands; Oceania includes Australia and New Zealand. No declines have been reported in Asia. [Photo credits (left to right): *Anaxyrus boreas*, Chris Brown; *Telmatobius sanborni*, Ignacio De la Riva; *Atelopus varius*, Brian Gratwicke; *Cycloramphus boraceiensis*, L. F. Toledo; *Cardioglossa melanogaster*, Mareike Hirschfeld; *Salamandra salamandra*, Didier Descouens; *Pseudophryne corroboree*, Corey Doughty].

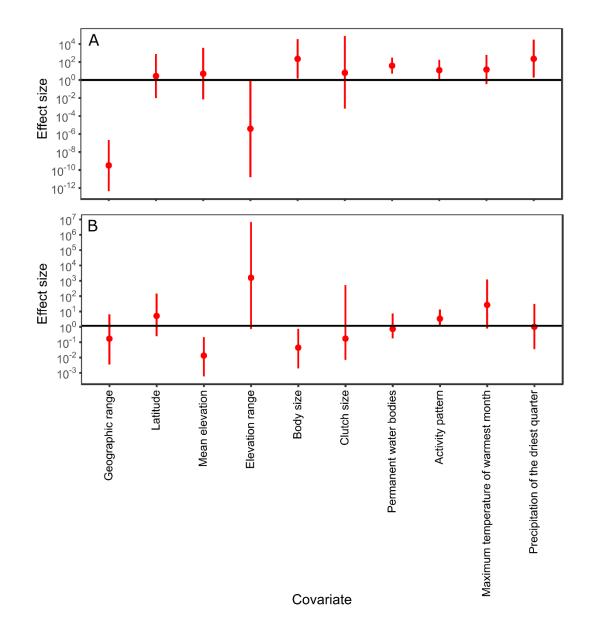


Figure S2. Multiplicative effect sizes for predictors of (A) decline severity and (B) recovery. Values greater and smaller than one indicate positive and negative effects respectively; a value of one (horizontal line) suggests no effect. Values are calculated from the ranges and coefficients in tables S2 and S3. Bars indicate 95% credible intervals.



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Table S1. Summary of modelling results for the timing of species declines. Estimates refer to the mean estimate of the regression coefficients for each model, respectively, intercept α , linear term β_{time} and quadratic term β_{time2} (95% credible intervals in parentheses). "Retained" refers to whether the variable was retained as significantly influencing the severity of declines (when the 95% credible interval of the regression coefficient's posterior distribution did not encompass zero).

Model	<u>Mean β (95% CRI)</u>	Retained
Total	number of declines (n=449)	
Decline ~ .	$\alpha = 2.25 \ (2.16, 2.34)$	No
Decline ~ time	$\alpha = 2.53 \ (2.35, 2.71)$	No
	$\beta_{\text{time}} = -1.13 (-1.84, -0.41)$	
Decline ~ time ²	$\alpha = 1.28 \ (0.94, \ 1.59)$	Yes
	$\beta_{time} = 14.37 \ (11.33, \ 17.51)$	
	$\beta_{time2} = -34.88 \ (-41.86, \ 28.19)$	
Se	everity of decline (n=449)	
Severity ~ .	α = -1.46 (-7.08, 3.79)	No
Severity ~ time	$\alpha = -1.32 (-6.08, 4.02)$	Yes
	$\beta_{time} = -9.65 (-15.36, -3.96)$	
Severity ~ time ²	$\alpha = -0.17 (-6.04, 4.75)$	No
	$\beta_{\text{time}} = -9.04 \ (-18.37, \ 0.31)$	
	$\beta_{time2} = -0.84 (-17.46, 15.68)$	
	Recovery (n=254)	1



Recovery ~ .	$\alpha = -1.86 (-3.05, -0.92)$	No
Recovery ~ time	$\alpha = -0.86 (-2.15, 0.33)$	Yes
	$\beta_{time} = -5.45 (-10.14, -1.04)$	
Recovery ~ time ²	$\alpha = -1.13 (-2.52, 0.15)$	No
	$\beta_{\text{time}} = -1.85 \ (-10.12, \ 6.19)$	
	$\beta_{\text{time2}} = -8.49 \ (-24.56, 7.55)$	



Variable

Latitude

Geographic range (km²)

Mean elevation (m asl)

Body size (SVL in mm)

Permanent water bodies

(strictly nocturnal yes/no)

Elevation range (m)

Clutch size (n)

Activity pattern

(yes/no)

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Table S2. Summary of results for predictors of the severity of species declines. "Range" refers to the range of a variable in the largest analysed dataset (all continuous variables were centred and modelled on the log_{10} scale). " β " refers to the mean estimate of the regression coefficient for each variable (95% credible intervals in parentheses). "Sample size" refers to the size of the largest fitted dataset which included the variable. "Retained" refers to whether the variable was retained as significantly influencing the severity of declines (when the 95% credible interval of the regression coefficient's posterior distribution did not encompass zero).

Mean β (95% CRI)

-2.60 (-3.39, -1.83)

0.46 (-2.12, 3.01)

0.87 (-2.69, 4.46)

-2.25 (-4.49, -0.03)

4.09 (0.28, 7.93)

0.53 (-2.10, 3.25)

3.66 (1.64, 5.72)

2.5 (-0.26, 5.15)

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Range (log10)

-5.04, 3.36

-1.58, 0.62

-1.24, 0.61

-4.84, 0.70

-0.55, 0.77

-1.50, 1.97

0, 1

0, 1

Retained

Yes

No

No

Yes

Yes

No

Yes

No

Sample size

297

217

217

297

297

102

297

217



Maximum temperature of	-0.27, 0.15	6.35 (-2.55, 15.26)	297	No
warmest month (C)				
Precipitation of the driest	-1.26, 0.88	2.53 (0.28, 4.81)	297	Yes
quarter (mm)				



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Table S3. Summary of results for predictors of species recovery (yes/no). "Range" refers to the range of a variable in the largest analysed dataset (all continuous variables were centred and modelled on the \log_{10} scale). " β " refers to the regression coefficient for the variable (on the \log_{10} scale for continuous variables). "Sample size" refers to the size of the largest fitted dataset which included the variable (number of known recoveries in parentheses). "Retained" refers to whether the variable was retained as significantly influencing the probability of recovery.

Variable	Range	Mean β (95% CRI)	Sample size	Retained
Geographic range (km ²)	-11.86, 7.47	-0.10 (-0.30, 0.09)	209 (50)	No
Latitude	-1.60, 0.62	0.67 (-0.70, 2.18)	210 (48)	No
Mean elevation (m asl)	-1.06, 0.49	-2.59 (-4.82, -0.53)	164 (41)	Yes
Elevation range (m)	-2.89, 0.83	1.94 (-0.13, 4.19)	164 (41)	No
Body size (SVL in mm)	-0.46, 0.77	-2.92 (-5.39, -0.67)	164 (41)	Yes
Clutch size (n)	-1.52, 1.85	-0.57 (-1.52,1.81)	84 (28)	No
Permanent water bodies	0, 1	-0.47 (-0.87, 1.87)	164 (41)	No
(yes/no)				
Activity pattern	0, 1	1.07 (0.05, 2.46)	154 (37)	Yes
(strictly nocturnal yes/no)				
Maximum temperature of	-0.27, 0.15	7.66 (-1.01, 16.98)	158 (40)	No
warmest month (C)				

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Precipitation of the driest	-1.30, 0.87	-0.07 (-1.61, 1.52)	158 (40)	No
quarter (mm)				



Table S4. Complete legend of taxonomic groups as indicated in Figure 2 (main text).

) rder	Family	Genus	Species	Severity	Recovery
nura	Alsodidae	Alsodes	Alsodes tumultuosus	1	no
	Alytidae	Alytes	Alytes muletensis	1	yes
			Alytes obstetricans	1	unknown
		Discoglossus	Discoglossus sardus	1	unknown
	Aromobatidae	Allobates	Allobates alessandroi	3	no
			Allobates capixaba*	3	yes
			Allobates olfersioides	4	n/a
			Allobates talamancae	2	unknown
		Aromobates	Aromobates meridensis	3	no
		Mannophryne	Mannophryne collaris	3	no
			Mannophryne cordilleriana	3	no
	Arthroleptidae	Arthroleptis	Arthroleptis variabilis	1	no
		Cardioglossa	Cardioglossa manengouba	3	no
			Cardioglossa pulchra	2	no no
			Cardioglossa trifasciata	3	no
	Brachycephalidae	Brachycephalus	Brachycephalus alipioi	2	no no
		Ischnocnema	Ischnocnema epipeda	3	no
			Ischnocnema paranaensis	4	n/a
			Ischnocnema parva	1	yes
			Ischnocnema pusilla	3	no
			Ischnocnema sp. aff. guentheri	1	yes
	Bufonidae	Anaxyrus	Anaxyrus baxteri	5	n/a
			Anaxyrus boreas	2	2 no
			Anaxyrus canorus	2	no no
		Atelopus	Atelopus andinus	2	unknown
			Atelopus angelito	4	n/a
			Atelopus arthuri	4	n/a
			Atelopus balios	3	no
			Atelopus bomolochos	3	no
			Atelopus boulengeri	4	n/a
			Atelopus carbonerensis	4	n/a
			Atelopus certus	3	no
			Atelopus chiriquiensis	4	n/a
			Atelopus chrysocorallus	4	n/a
			Atelopus coynei	4	n/a
			Atelopus cruciger	3	unknown
			Atelopus dimorphus	4	n/a
			Atelopus elegans	3	no
			Atelopus epikeisthos	4	n/a



Order Family	Genus	Species	Severity	Recovery
		Atelopus eusebiodiazi		4 n/a
		Atelopus exiguus		<mark>3</mark> no
		Atelopus glyphus		<mark>3</mark> no
		Atelopus guanujo		4 n/a
		Atelopus halihelos		4 n/a
		Atelopus ignescens		<mark>3</mark> no
		Atelopus limosus		<mark>3</mark> no
		Atelopus longirostris		<mark>3</mark> no
		Atelopus lynchi		4 n/a
		Atelopus mindoensis		4 n/a
		Atelopus mucubajiensis		<mark>3</mark> no
		Atelopus nanay		<mark>3</mark> no
		Atelopus nepiozomus		<mark>3</mark> no
		Atelopus onorei		4 n/a
		Atelopus orcesi		4 n/a
		Atelopus oxyrhynchus		4 n/a
		Atelopus pachydermus		<mark>3</mark> no
		Atelopus palmatus		<mark>3</mark> no
		Atelopus pastuso		4 n/a
		Atelopus patazensis		<mark>3</mark> yes
		Atelopus peruensis		4 n/a
		Atelopus petersi		4 n/a
		Atelopus pinangoi		4 n/a
		Atelopus planispina		4 n/a
		Atelopus podocarpus		4 n/a
		Atelopus pulcher		<mark>3</mark> no
		Atelopus pyrodactylus		4 n/a
		Atelopus reticulatus		4 n/a
		Atelopus senex		4 n/a
		Atelopus sorianoi		4 n/a
		Atelopus sp.		4 n/a
		Atelopus spurrelli		3 unknown
		Atelopus tamaense		4 n/a
		Atelopus tricolor		<mark>3</mark> no
		Atelopus varius		<mark>3</mark> yes
		Atelopus zeteki		4 n/a
	Incilius	Incilius aucoinae		1 unknown
		Incilius coniferus		1 unknown
		Incilius epioticus		1 unknown
		Incilius fastidiosus		$\frac{1}{3}$ no
		Incilius holdridgei		3 no
		Incilius ibarrai		3 no



rder	Family	Genus	Species	Severity	Recovery
			Incilius macrocristatus	2	no
			Incilius periglenes	5	n/a
			Incilius signifer	1	unknown
			Melanophryniscus moreirae	3	yes
		Nectophrynoides	Nectophrynoides asperginis	5	n/a
		Rhaebo	Rhaebo haematiticus	2	yes
		Rhinella	Rhinella alata	1	unknown
			Rhinella arunco	1	no
			Rhinella centralis	1	unknown
			Rhinella leptoscelis	3	yes
			Rhinella quechua	3	no
			Rhinella spinulosa	2	yes
			Rhinella veraguensis	3	no
		Werneria	Werneria tandyi	2	no
	Calyptocephalellidae	Calyptocephalella	Calyptocephalella gayi	2	no
	Centrolenidae	Centrolene	Centrolene buckleyi	1	unknown
			Centrolene daidaleum	1	no
		Cochranella	Cochranella euknemos	2	no
			Cochranella granulosa	1	no
		Espadarana	Espadarana prosoblepon	1	unknown
		Hyalinobatrachium	Hyalinobatrachium aureoguttatum	2	unknown
			Hyalinobatrachium bergeri	1	no
			Hyalinobatrachium chirripoi	1	unknown
			Hyalinobatrachium colymbiphyllum	2	yes
			Hyalinobatrachium fleischmanni	1	yes
			Hyalinobatrachium talamancae	2	unknown
		Hyalinobatrachium	Hyalinobatrachium valerioi	1	unknown
		Nymphargus	Nymphargus griffithsi	1	no
			Nymphargus pluvialis	3	no
			Nymphargus truebae	4	n/a
		Rulyrana	Rulyrana spiculata	3	no
		Sachatamia	Sachatamia albomaculata	2	yes
			Sachatamia ilex		unknown
		Teratohyla	Teratohyla pulverata	1	unknown
			Teratohyla spinosa		unknown
		Vitreorana	Vitreorana eurygnatha		yes
	Craugastoridae	Bryophryne	Bryophryne cophites		no
			Bryophryne zonalis		unknown
		Craugastor	Craugastor anciano		n/a
			Craugastor andi		no
			Craugastor angelicus		no



Order Family	Genus	Species	Severity	Recovery
		Craugastor aurilegulus		<mark>3</mark> no
		Craugastor azueroensis		<mark>3</mark> unknown
		Craugastor bransfordii		2 unknown
		Craugastor catalinae		<mark>3</mark> no
		Craugastor chrysozetetes		5 n/a
		Craugastor crassidigitus		2 unknown
		Craugastor cruzi		4 n/a
		Craugastor emleni		<mark>4</mark> n/a
		Craugastor escoces		<mark>3</mark> unknown
		Craugastor evanesco		<mark>3</mark> no
		Craugastor fitzingeri		2 unknown
		Craugastor fleischmanni		<mark>3</mark> no
		Craugastor gollmeri		<mark>3</mark> unknown
		Craugastor megacephalus		2 yes
		Craugastor melanostictus		1 unknown
		Craugastor mexicanus		2 no
		Craugastor milesi		<mark>3</mark> no
		Craugastor monnichorum		2 unknown
		Craugastor noblei		<mark>3</mark> unknown
		Craugastor obesus		4 n/a
		Craugastor omoaensis		4 n/a
		Craugastor opimus		2 unknown
		Craugastor podiciferus		1 unknown
		Craugastor punctariolus		4 n/a
		Craugastor ranoides		<mark>3</mark> no
		Craugastor rhyacobatrachus		4 n/a
		Craugastor rugosus		2 unknown
		Craugastor stadelmani		4 n/a
		Craugastor stejnegerianus		1 unknown
		Craugastor tabasarae		<mark>3</mark> no
		Craugastor talamancae		2 unknown
		Craugastor taurus		<mark>3</mark> no
		Craugastor underwoodi		2 unknown
	Holoaden	Holoaden bradei		4 n/a
	Pristimantis	Pristimantis anolirex		2 no
		Pristimantis caryophyllaceus		2 yes
		Pristimantis cerasinus		2 unknown
		Pristimantis cosnipatae		3 unknown
		Pristimantis cruentus		2 unknown
		Pristimantis danae		1 unknown
		Pristimantis gaigei		2 yes
		Pristimantis gracilis		1 no



Order	Family	Genus	Species	Severity	Recovery
			Pristimantis gryllus	1	no
			Pristimantis melanoproctus	2	2 no
			Pristimantis mondolfii	1	no
			Pristimantis moro	2	2 unknown
			Pristimantis museosus	3	unknown
			Pristimantis nicefori	1	no
			Pristimantis palmeri	1	yes
			Pristimantis pardalis	2	2 yes
			Pristimantis pharangobates	1	unknown
			Pristimantis platydactylus	1	yes
			Pristimantis ridens	2	2 yes
			Pristimantis salaputium	1	unknown
			Pristimantis taeniatus	1	unknown
			Pristimantis toftae]	yes
		Strabomantis	Strabomantis bufoniformis	3	no
		Tachiramantis	Tachiramantis douglasi	2	2 no
	Cycloramphidae	Cycloramphus	Cycloramphus boraceiensis	1	no
			Cycloramphus duseni	2	n/a
			Cycloramphus fuliginosus	1	no
			Cycloramphus granulosus	3	no
			Cycloramphus semipalmatus	2	2 no
			Cycloramphus stejnegeri	۷	n/a
			Cycloramphus valae	۷	n/a
		Thoropa	Thoropa petropolitana	3	no
			Thoropa taophora	1	no
	Dendrobatidae	Ameerega	Ameerega flavopicta	1	yes
		Andinobates	Andinobates minutus		unknown
		Colostethus	Colostethus panamansis		yes
			Colostethus pratti		no
		Dendrobates	Dendrobates auratus		2 yes
		Hyloxalus	Hyloxalus abditaurantius		no
		•	Hyloxalus fascianigrus		no no
			Hyloxalus lehmanni		no
		Oophaga	Oophaga arborea		unknown
		1 0			2 unknown 2 unknown
			Oophaga granulifera Oophaga histrionica		2 unknown 2 no
			Oophaga nisirionica Oophaga pumilio		unknown
					n/a
			Oophaga speciosa		
		Phyllobates	Oophaga vicentei		unknown
		Ranitomeya	Phyllobates lugubris		unknown
		Runtonicya	Ranitomeya claudiae	2	unknown



Order	Family	Genus	Species	Severity	Recovery
		Silverstoneia	Silverstoneia flotator		2 yes
			Silverstoneia nubicola	4	2 unknown
	Eleutherodactylidae		Adelophryne baturitensis		<mark>3</mark> no
		Diasporus	Diasporus diastema	-	lunknown
			Diasporus hylaeformis	-	lunknown
			Diasporus quidditus	-	unknown
		Eleutherodactylus	Eleutherodactylus coqui	-	l yes
			Eleutherodactylus eneidae	4	5 n/a
			Eleutherodactylus jasperi	4	5 n/a
			Eleutherodactylus karlschmidti	4	5 n/a
			Eleutherodactylus locustus	3	<mark>3</mark> yes
			Eleutherodactylus portoricensis		l yes
			Eleutherodactylus richmondi		<mark>3</mark> yes
			Eleutherodactylus wightmanae	4	2 yes
	Hemiphractidae	Fritziana	Fritziana ohausi	-	l no
		Gastrotheca	Gastrotheca antoniiochoai		unknown
			Gastrotheca cornuta		<mark>3</mark> no
			Gastrotheca dendronastes	-	l no
			Gastrotheca excubitor		no
			Gastrotheca helenae	1	2 no
			Gastrotheca nebulanastes	-	l no
			Gastrotheca nicefori	-	l no
			Gastrotheca testudinea	-	l no
		Hemiphractus	Hemiphractus fasciatus		2 unknown
			Hemiphractus helioi	-	l no
	Hylidae	Aplastodiscus	Aplastodiscus flumineus	2	1 n/a
			Aplastodiscus musicus		<mark>3</mark> no
		Boana	Boana boans	-	unknown
			Boana gladiator	-	unknown
			Boana pugnax	-	unknown
			Boana rosenbergi	-	lunknown
			Boana rufitela		2 unknown
			Boana xerophyla	-	l no
		Bokermannohyla	Bokermannohyla circumdata	-	l no
			Bokermannohyla izecksohni		<mark>3</mark> no
		Bromeliohyla	Bromeliohyla bromeliaceae		<mark>3</mark> no
		Charadrahyla	Charadrahyla altipotens		3 no
			Charadrahyla nephila		2 no
		Cruziohyla	Cruziohyla calcarifer		unknown
		Dendropsophus	Dendropsophus ebraccatus		unknown
			Dendropsophus meridensis		2 no



Order

Family	Genus	Species	Severity	Recovery
		Dendropsophus microcephalus	1	unknown
		Dendropsophus pelidna	1	no
		Dendropsophus phlebodes	1	unknown
		Dendropsophus ruschii	3	yes
	Diaglena	Diaglena spinosa	2	unknown
	Dryophytes	Dryophytes euphorbiacea	1	no
		Dryophytes walkeri	2	no
	Duellmanohyla	Duellmanohyla ignicolor	2	no
		Duellmanohyla schmidtorum	2	no
		Duellmanohyla soralia	2	no
		Duellmanohyla uranochroa	3	no
	Ecnomiohyla	Ecnomiohyla echinata	2	no
		Ecnomiohyla miliaria	2	no
		Ecnomiohyla minera	3	no
		Ecnomiohyla rabborum	4	n/a
	Exerodonta	Exerodonta melanomma	2	no
	Hyla	Hyla bocourti	4	n/a
		Hyla hazelae	2	unknown
	Hyloscirtus	Hyloscirtus armatus	3	no
		Hyloscirtus colymba	2	no
		Hyloscirtus palmeri	3	unknown
		Hyloscirtus phyllognathus	3	no
		Hyloscirtus platydactylus	1	no
	Isthmohyla	Isthmohyla angustilineata	3	no
		Isthmohyla calypsa	4	n/a
		Isthmohyla debilis	3	no
		Isthmohyla graceae	3	no
		Isthmohyla lancasteri	2	unknown
		Isthmohyla picadoi	1	unknown
		Isthmohyla pictipes	2	no
		Isthmohyla pseudopuma	2	no
		Isthmohyla rivularis	3	no
		Isthmohyla tica	3	no
		Isthmohyla zeteki	1	unknown
	Ololygon	Ololygon cf. perpusillus	2	yes
		Ololygon heyeri	3	yes
		Ololygon peixotoi	3	no
	Phyllodytes	Phyllodytes luteolus	1	no
	Plectrohyla	Plectrohyla cyclada	2	no
		Plectrohyla dasypus		no
		Plectrohyla exquisita		no



rder	Family	Genus	Species	Severity	Recovery
			Plectrohyla hartwegi		2 no
			Plectrohyla ixil		<mark>3</mark> no
			Plectrohyla lacertosa		<mark>3</mark> no
			Plectrohyla matudai		2 no
			Plectrohyla quecchi		2 no
			Plectrohyla sagorum		2 no
		Pseudis	Pseudis paradoxa		1 no
		Ptychohyla	Ptychohyla erythromma		2 no
			Ptychohyla hypomykter		1 yes
			Ptychohyla legleri		<mark>3</mark> no
			Ptychohyla leonhardschultzei		2 no
		Sarcohyla	Sarcohyla arborescandens		2 no
			Sarcohyla sabrina		<mark>3</mark> no
		Scarthyla	Scarthyla vigilans		1 no
		Scinax	Scinax altae		1 unknown
			Scinax boulengeri		1 unknown
			Scinax elaeochrous		1 unknown
			Scinax ruber		1 unknown
		Smilisca	Smilisca phaeota		2 yes
			Smilisca sila		1 yes
			Smilisca sordida		1 unknown
		Tepuihyla	Tepuihyla edelcae		1 no
		Tlalocohyla	Tlalocohyla loquax		2 no
		Trachycephalus	Trachycephalus typhonius		1 unknown
	Hylodidae	Crossodactylus	Crossodactylus cf. gaudichaudii		1 no
			Crossodactylus dispar		4 n/a
			Crossodactylus timbuhy		3 yes
			Crossodactylus trachystomus		3 yes
			Crossodactylus werneri		3 no
	Ī	Hylodes	Hylodes asper		1 no
			Hylodes babax		1 no
			Hylodes glaber		4 n/a
			Hylodes lateristrigatus		1 no
			Hylodes phyllodes		1 yes
	Hyperoliidae	Afrixalus	Afrixalus paradorsalis		1 no
		Kassina	Kassina decorata		2 no
	Leiopelmatidae	Leiopelma	Leiopelma archeyi		1 unknown
	Leptodactylidae	Adenomera	Adenomera andreae		1 no
	- •		Adenomera marmorata		1 yes
		Leptodactylus	Leptodactylus bolivianus		1 unknown
			Leptodactylus colombiensis		1 no



er	Family	Genus	Species	Severity	Recovery
			Leptodactylus fallax	3	<mark>8</mark> no
			Leptodactylus fragilis	1	unknown
			Leptodactylus melanonotus	1	unknown
			Leptodactylus rhodonotus	1	no
			Leptodactylus savagei	1	unknown
			Leptodactylus sp.	3	no no
		Paratelmatobius	Paratelmatobius lutzii	4	n/a
		Physalaemus	Physalaemus henselii	2	2 no
			Physalaemus jordanensis	2	2 no
		Pleurodema	Pleurodema bibroni	2	2 no
			Pleurodema brachyops	1	unknown
			Pleurodema marmoratum]	unknown
	Limnodynastidae	Adelotus	Adelotus brevis	1	no
	Microhylidae	Chiasmocleis	Chiasmocleis panamensis	1	unknown
		Ctenophryne	Ctenophryne aterrima	1	unknown
		Hypopachus	Hypopachus barberi	2	2 no
	Myobatrachidae	Geocrinia	Geocrinia victoriana	1	no
		Mixophyes	Mixophyes balbus	2	yes
			Mixophyes fleayi		yes
			Mixophyes iteratus		yes
		Philoria	Philoria frosti		no
		Pseudophryne	Pseudophryne bibronii	1	no
			Pseudophryne corroboree	3	no
			Pseudophryne dendyi	1	no
			Pseudophryne pengilleyi	3	no
		Rheobatrachus	Rheobatrachus silus	4	n/a
			Rheobatrachus vitellinus	4	n/a
		Taudactylus	Taudactylus acutirostris	4	5 n/a
			Taudactylus diurnus	4	5 n∕a
			Taudactylus eungellensis	2	2 yes
			Taudactylus liemi		yes
			Taudactylus pleione		<mark>8</mark> no
			Taudactylus rheophilus	4	n/a
	Odontophrynidae	Proceratophrys	Proceratophrys moratoi	1	no
	Pelodryadidae	Litoria	Litoria castanea	3	<mark>8</mark> no
			Litoria littlejohni	1	no
			Litoria verreauxii	1	yes
			Litoria verreauxii alpina	2	2 no
		Ranoidea	Ranoidea aurea	2	2 no
			Ranoidea barringtonensis	1	unknown
			Ranoidea booroolongensis	2	2 no



er	Family	Genus	Species	Severity	Recovery
			Ranoidea caerulea	1	no
			Ranoidea daviesae	1	unknown
			Ranoidea dayi	2	no
			Ranoidea jungguy	1	unknown
			Ranoidea kroombitensis	1	no
			Ranoidea lesueuri	1	unknown
			Ranoidea lorica	3	no
			Ranoidea nannotis	2	yes
			Ranoidea nudidigita	1	yes
			Ranoidea nyakalensis	5	n/a
			Ranoidea pearsoniana	2	yes
			Ranoidea piperata	4	n/a
			Ranoidea raniformis	2	no
			Ranoidea rheocola	2	yes
			Ranoidea serrata	1	yes
			Ranoidea spenceri	3	no
			Ranoidea subglandulosa	1	unknown
			Ranoidea wilcoxii	1	unknown
	Petropedetidae	Petropedetes	Petropedetes perreti	1	no
	Phrynobatrachidae	Phrynobatrachus	Phrynobatrachus cricogaster	1	no
			Phrynobatrachus jimzimkusi	2	no
			Phrynobatrachus manengoubensis	3	no
			Phrynobatrachus sandersoni	2	no
			Phrynobatrachus werneri	2	no
	Phyllomedusidae	Agalychnis	Agalychnis annae	3	no
			Agalychnis callidryas	1	unknown
			Agalychnis lemur	3	no
			Agalychnis moreleti	3	no
			Agalychnis spurrelli	1	unknown
		Phasmahyla	Phasmahyla exilis	1	no
			Phasmahyla guttata	1	yes
		Phrynomedusa	Phrynomedusa bokermanni	4	n/a
			Phrynomedusa marginata	4	n/a
		Phyllomedusa	Phyllomedusa venusta	1	unknown
	Pipidae	Pipa	Pipa myersi	2	unknown
	Ranidae	Lithobates	Lithobates chiricahuensis	2	no
			Lithobates maculatus	2	yes
			Lithobates megapoda	2	no
			Lithobates montezumae	2	no
			Lithobates neovolcanicus	1	no
			Lithobates pipiens	2	no



)rder	Family	Genus	Species	Severity	Recovery
			Lithobates sierramadrensis		2 no
			Lithobates spectabilis	:	2 no
			Lithobates tarahumarae		2 no
			Lithobates vaillanti		1 unknown
			Lithobates vibicarius		<mark>3</mark> yes
			Lithobates warszewitschii	:	2 yes
			Lithobates yavapaiensis		2 yes
		Rana	Rana cascadae		2 no
			Rana maculata		1 yes
			Rana muscosa		3 no
			Rana sierrae		3 yes
	Rhinodermatidae	Rhinoderma	Rhinoderma darwinii		2 no
			Rhinoderma rufum		4 n/a
	Telmatobiidae	Telmatobius	Telmatobius arequipensis		2 no
			Telmatobius atacamensis		4 n/a
			Telmatobius atahualpai		1 unknown
			Telmatobius bolivianus		4 n/a
			Telmatobius brevipes		2 no
			Telmatobius brevirostris		2 no
			Telmatobius carrillae		2 no
			Telmatobius ceiorum		4 n/a
			Telmatobius chusmisensis		1 no
			Telmatobius cirrhacelis		4 n/a
			Telmatobius colanensis		2 no
			Telmatobius contrerasi		4 n/a
			Telmatobius culeus		unknown
			Telmatobius edaphonastes		4 n/a
			Telmatobius espadai		4 n/a
			Telmatobius hockingi		2 no
			Telmatobius ignavus		2 no
			Telmatobius jelskii		1 unknown
			Telmatobius laticeps		4 n/a
			Telmatobius latirostris		2 unknown
			Telmatobius marmoratus		2 yes
			Telmatobius mayoloi		2 unknown
			Telmatobius mendelsoni		4 n/a
			Telmatobius niger		4 n/a
			Telmatobius pefauri		4 n/a
			Telmatobius pisanoi		4 n/a
			Telmatobius punctatus		1 no
			Telmatobius sanborni		2 no



Order	Family	Genus	Species	Severity	Recovery
			Telmatobius schreiteri		4 n/a
			Telmatobius scrocchii		4 n/a
			Telmatobius sibiricus		<mark>4</mark> n/a
			Telmatobius simonsi		<mark>3</mark> unknown
			Telmatobius stephani		<mark>4</mark> n/a
			Telmatobius timens		<mark>3</mark> no
			Telmatobius truebae		<mark>3</mark> no
			Telmatobius vellardi		2 no
			Telmatobius verrucosus		4 n/a
			Telmatobius yuracare		<mark>3</mark> n/a
laudata	Ambystomatidae	Ambystoma	Ambystoma altamirani		1 no
			Ambystoma granulosum		1 no
	Plethodontidae	Bolitoglossa	Bolitoglossa biseriata		1 unknown
			Bolitoglossa colonnea		2 unknown
			Bolitoglossa compacta		1 unknown
			Bolitoglossa leandrae		2 no
			Bolitoglossa lignicolor		1 unknown
			Bolitoglossa magnifica		1 unknown
			Bolitoglossa marmorea		1 unknown
			Bolitoglossa medemi		1 unknown
			Bolitoglossa minutula		1 unknown
			Bolitoglossa rostrata		2 no
			Bolitoglossa schizodactyla		<mark>3</mark> unknown
			Bolitoglossa tamaense		1 no
		Chiropterotriton	Chiropterotriton dimidiatus		2 no
			Chiropterotriton magnipes		<mark>3</mark> no
		Oedipina	Oedipina grandis		2 unknown
			Oedipina parvipes		1 unknown
		Parvimolge	Parvimolge townsendi		2 no
		Pseudoeurycea	Pseudoeurycea brunnata		4 n/a
			Pseudoeurycea cochranae		<mark>3</mark> no
			Pseudoeurycea exspectata		4 n/a
			Pseudoeurycea goebeli		<mark>3</mark> no
			Pseudoeurycea lineola		2 no
			Pseudoeurycea melanomolga		2 no
			Pseudoeurycea nigromaculata		2 no
			Pseudoeurycea papenfussi		<mark>3</mark> no
			Pseudoeurycea rex		2 no
			Pseudoeurycea smithi		<mark>3</mark> no
			Pseudoeurycea unguidentis		4 n/a



Order	Family	Genus	Species	Severity	Recovery
		Thorius	Thorius aureus		l no
			Thorius boreas		1 no
			Thorius dubitis		<mark>3</mark> no
			Thorius magnipes		<mark>3</mark> no
			Thorius pennatulus	2	2 no
			Thorius pulmonaris		2 no
			Thorius troglodytes		<mark>3</mark> no
	Salamandridae	Euproctus	Euproctus platycephalus		l unknown
		Salamandra	Salamandra salamandra		l no



Data S1. For the complete dataset of declined species, see the separate Excel file.