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Null expectations for disease dynamics in shrinking habitat: dilution or amplification?

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Main Text

Summary

As biodiversity declines with anthropogenic land-use change, it is increasingly important to understand how changing biodiversity affects infectious disease risk. The dilution effect hypothesis, which points to decreases in biodiversity as critical to an increase in infection risk, has received considerable attention due to the allure of a win-win scenario for conservation and human well-being. Yet some empirical data suggests that the dilution effect is not a generalizable phenomenon. We explore the response of pathogen transmission dynamics to changes in biodiversity that are driven by habitat loss using an allometrically-scaled multi-host model. With this model, we show that declining habitat, and thus declining biodiversity, can lead to either increasing or decreasing infectious disease risk, measured as endemic prevalence. Whether larger habitats, and thus greater biodiversity, lead to a decrease (dilution effect) or increase (amplification effect) in infection prevalence depends upon the pathogen transmission mode and how host competence scales with body size. Dilution effects were detected for most frequency-transmitted pathogens and amplification effects were detected for most density-dependent pathogens. Amplification effects were also observed over a particular range of habitat loss in frequency dependent pathogens when we assumed that host competence was greatest in large bodied species. By contrast, only amplification effects were observed for density dependent pathogens; host competency only affected the magnitude of the effect. These models can be used to guide future empirical studies of biodiversity-disease relationships across gradients of habitat loss. The type of

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transmission, the relationship between host competence and community assembly, the identity of hosts contributing to transmission, and how transmission scales with area, are will be essential factors to consider when elucidating the mechanisms driving disease risk in shrinking habitat.

Introduction

Understanding how habitat loss affects pathogen prevalence and infection risk is a critical challenge in the face of unprecedented rates of anthropogenic landscape transformation [1]. It is clear that habitat loss leads to biodiversity loss [2-4], but it remains uncertain how declining habitat and diversity affect the transmission of infectious diseases [5]. A dominant hypothesis, known as the dilution effect, states that declining biodiversity leads to increased infectious disease transmission [6, 7]. The rationale is that greater host diversity provides a higher proportion of low competent hosts and therefore 'dilutes' the transmission chain. This creates a strong incentive to conserve intact ecological communities from the perspective of human health.

Despite the appeal of applying the dilution effect to guide win-win management strategies for conservation and public health, the empirical evidence has been inconsistent. Some studies suggest that increasing host diversity decreases infectious disease risk (a 'dilution effect') [7-12], while others suggest increasing diversity increases infectious disease risk (an amplification effect) [9, 13]. Large-scale surveys of West Nile Virus and Lyme disease suggested that communities with higher diversity (measured as species richness) have a lower prevalence of disease [14, 15]. By contrast, surveys of trophically-transmitted (pathogens transmitted through food webs) and host-specific pathogens, have suggested that intact ecosystems have higher pathogen prevalence than less-diverse system. For example, losses of host populations due to fragmentation in Californian estuarine ecosystems led to a reduction in observed pathogen prevalence [16-19]. These conflicting patterns in empirical data have resulted in considerable academic debate about the generalizability of the dilution effect [9, 20-24].

An increasing number of studies have investigated relationships between biodiversity and disease risk and subsequent meta-analyses have attempted to identify consistent patterns. The majority of empirical studies use prevalence as a proxy for infection risk. By comparing communities with one host species to communities with at least two host species, Civitello and colleagues [25] found support for lower disease risk in multi-species communities compared with single species populations across a variety of pathogen systems. In contrast, Salkeld and colleagues [23] found minimal support for the dilution effect amongst zoonotic pathogens. Salkeld et al. also highlighted a potential publication bias for studies supporting the dilution effect [23] - small studies that do not support the dilution effect have not been published. The allure of the dilution effect may be influencing the publication rates and overselling it as a generalizable phenomena.

In parallel to empirical studies, models of disease transmission in multi-host parasite systems have been developed to understand when species diversity and community structure impact parasite transmission [26-29]. Theory suggests that the type of pathogen transmission determines the outcome of diversity-disease relationships. Multi-host pathogens with frequency dependent transmission are expected to decrease in prevalence as biodiversity increases, i.e. the dilution effect [28, 29]. Vector-borne pathogen transmission is typically frequency-dependent and dilution effects are predicted [28, 29], however, vector preference and variation in host competence can complicate the outcome of these diversity-disease relationships [30-32]. Pathogens that follow density-dependent transmission are expected to increase as host diversity increases [28]. While amplification effects are predicted for most density-dependent pathogens, when total abundance saturates well before species richness, both amplification and dilution effects can be observed [33].

Most theoretical studies of disease and diversity rely on measuring changes in community R_0 . Community R_0 describes the number of secondary cases caused by the introduction of a single infectious individual into a completely naïve community of multiple host species. Community R_0 is a proxy for infection risk - it indicates whether or not an epidemic will occur ($R_0 \geq 1$) and is linked to peak prevalence of an epidemic. In simple cases R_0 can be used to infer endemic equilibrium, but numerical simulations are required to understand more complex multi-host pathogens (Keeling and Rohani). Methods have been developed to use equilibrium prevalence of a disease to infer community R_0 (see [34]), but oftentimes there is a disconnect between models predictions and available empirical data.

While habitat loss is a major driver of declining biodiversity [35], this process has not yet been linked explicitly to models of disease. Landscapes undergoing habitat loss often have lower biodiversity, both in number of species and evenness of communities. Empirical studies of mammalian disease burdens often report higher parasite prevalence in fragmented landscapes [36-44]. By contrast, a positive relationship between parasite prevalence and patch size has been observed in several wild bird populations [45-47]. Relationships between habitat patch size, host diversity, and pathogen transmission within a habitat remain relatively unexplored (although see [5]). While most studies of parasites across a gradient of habitat loss do not directly measure biodiversity, they offer the opportunity to understand how real-world processes affecting community composition impact disease transmission.

Here, we develop null expectations for disease transmission in a single habitat patch undergoing loss of area and host diversity. We explore how a directly-transmitted pathogen responds to changing habitat size in a multi-host community. Specifically, we ask: 1) How does habitat patch size affect the dynamics and prevalence of a generalist pathogen with density-dependent transmission? 2) How does patch size affect a generalist pathogen with frequency-dependent transmission? 3) How does the order of community disassembly affect these results? 4) What assumptions about host competence are necessary to observe dilution versus amplification effects in an area undergoing habitat loss? and 5) How does changing assumptions of host range affect transmission of density dependent pathogens?

Methods

Definitions of biodiversity and disease risk. An array of terminology and measurements are used in the dilution and amplification effect literature to describe the relationships between host biodiversity and disease (reviewed by [9, 24]). To keep the results of simulations consistent with empirical observations, we used species richness and equilibrium disease prevalence to measure biodiversity and disease, respectively. Unless indicated otherwise, we used the terminology and metrics interchangeably.

Model Assumptions and Formulation.

Disease free equilibrium. We examined community composition and resultant disease dynamics across a gradient of habitat size, from 0.1 km^2 (10 hectares) to 100 km^2 (10,000 hectares). We assumed there were up to ten host species that contributed to pathogen transmission in these simulations. The average body mass (m) of individuals from each of the ten host species was randomly selected from a skewed normal distribution fit to observed distributions of terrestrial mammalian body sizes [48, 49] (Fig. S1, ESM Methods).

The carrying capacity (K_{ia}) for each species (i) was determined by its body-mass determined density ($16.2m_i^{-0.70}$ per km^2) and a given habitat size (a). These assumptions gave us reasonable species accumulation curves at disease free equilibrium: increasing habitat size increased the number of individuals in a given patch and the number of species, realistically mirroring patterns observed in natural systems (Fig. S2-S4). We did not take into account interspecies interactions such as competition, as other authors have explored the impacts of these interactions on disease dynamics [50].

The community of hosts at a given habitat size was determined following one of three community assembly rules: (1) a species was resident in a patch if the carrying capacity was above a fixed threshold of 10 individuals, (2) a species was considered resident if the carrying capacity was above the species-specific threshold which was randomly selected from a uniform distribution (1-100), or (3) a unique habitat density modifier (ϵ_{ia}) was randomly selected from a uniform distribution (0,2) for each species (i), and thus density of each host species varies at each habitat area. In the first two scenarios, species are perfectly nested – i.e. once they are extinct in a habitat patch, they never recolonize as the habitat continues to shrink. This constant density across habitat size is supported by empirical data [26]. The second community assembly rule is akin to some species being more tolerant to habitat loss (low population threshold) compared with other species that may be highly sensitive (high population threshold). The third assembly rule allows variation in species responses to habitat size-- this can be due to variations in resource requirements, sensitivity to edge effects, or another confounding factor.

Disease model and assumptions. The deterministic multi-host model was modified from an allometrically-scaled S-I-R model for microparasites [51]. For each species i , susceptible (S_i), infected (I_i), and recovered (R_i) numbers of hosts were modeled (density-dependent transmission shown in eqs. 1-3, frequency-dependent transmission in eqs S1-S3):

$$\frac{dS_i}{dt} = b_i N_i \left(1 - \frac{N_i}{K_{ia}}\right) - \frac{S_i(\beta_{ii}I_i + \psi \sum_j^n \beta_{ij}I_j)}{a} - d_i S_i \quad (1)$$

$$\frac{dI_i}{dt} = \frac{S_i(\beta_{ii}I_i + \psi \sum_j^n \beta_{ij}I_j)}{a} - I_i(\alpha_i + \gamma_i + d_i) \quad (2)$$

$$\frac{dR_i}{dt} = \gamma_i I_i - d_i R_i \quad (3)$$

In these models, we are assuming microparasites are transmitted through direct contact with conspecifics and heterospecifics. For density dependent pathogens, transmission rates are dependent on the baseline force of infection and the density of infected and uninfected hosts (SI/a , where a is the total area of the patch). In contrast, frequency dependent pathogen transmission is related to the proportion of infected individuals in the population (SI/N , where N is the total number of individuals in a patch). Conceptualizing transmission in this way focuses on two ends of what is likely a continuum [52], but it is a convenient way to understand the spectrum of transmission modes.

Demographic and disease parameters were determined by allometric scaling (following [51] detailed in Table S1). For baseline models, it was assumed that R_0 within each species was constant (eqs. S4, S5) - an isolated, closed, non-breeding population of each species would experience similar disease epidemics and host competence did not vary between species. R_0 is often thought of as a static parameter, but it is dependent on the population into which it is introduced. R_0 for a single species is different than the community R_0 . For multihost pathogens, the number of secondary cases depend not only on characteristics of a single host species, but of all species capable of transmission in the community. In the methods, we will use R_0 to talk mainly about the species-specific parameter, but will specify when we are referring to R_0 for the whole community.

Within species transmission rates (β_{ii}) were calculated using species-specific recovery, mortality, and disease induced mortality rates (Table S1). We assumed that between species transmission rates were the product of the between species transmission rate (β_{ij}), which was the geometric mean of the donor (i) and recipient (j) within species transmission rates ($\sqrt{\beta_{ii} \beta_{jj}}$), and a between species contact rate, which was a proportion of within species contact (ψ). The between species contact allowed us to vary interactions between species from zero (non-overlapping contact patterns; $\psi = 0$), to one (between species contact was nearly equivalent to within-species contacts; $\psi = 1$). Simulations were run to equilibrium at a given patch size ($t = 150$ years) and resultant prevalence in each species and across the community were calculated. In the above model set up, we assumed that host competence, defined as within species R_0 (Eq. S4, S5), was the same for each species.

Variation in host competence. An alternative scenario is that host competence varies between species, and therefore within species R_0 is not constant across body size for a given pathogen. Some authors argue that life history trade-offs dictate a negative relationship between R_0 and body size [54], meaning smaller species are more competent hosts. Species with faster life histories, indicating smaller body sizes, have been shown to acquire and transmit infections better than slow-lived species [55, 56]. This variation in competence could be related to lower investment in adaptive immunity in fast-living species [57, 58]. We scaled β_{ii} as a function of body size, ensuring that the smallest bodied species' R_0 was the largest (Fig S5).

Conversely, if behavioral allometry is taken into account, R_0 is expected to increase with body size [59]. This increase in R_0 is mediated by increasing group size and social contacts in larger bodied animals and is supported by prevalence data for ungulates and primates from the Global Mammalian Database [59]. In some simulations, we included conditions of behavioral allometry, scaling β_{ii} so that the largest R_0 was found in the largest host species (Figure S5).

Lastly, we examined the effect of variation in host competence by assembling a community that included incompetent hosts unrelated to body size (and thus unrelated to community assemblage in this model). Incompetent hosts could get infected but could not transmit the pathogen onwards ($\beta_{ii} = \beta_{ij} = 0$; within and between incompetent host transmission rates were set to zero).

Variation in evenly-mixed populations. Finally, we considered a null model of community pathogen dynamics with allometric scaling of home ranges. In the baseline model we considered infection risk of a density dependent pathogen to scale with individuals per km^2 . However, individual species may not come into contact with all individuals in a square kilometer. To account for heterogeneity in between species contact, we used approximate home ranges as the area in which contacts occur. The minimum within species transmission rate, $\beta_{ii} = 0.25 \left(md_i + \frac{y_i}{d_i} \right) m_i^{-0.69}$ - is similar to density dependent transmission calculated previously [51] but includes the assumption that home range scales with body mass. In addition to a different transmission parameter, the system of ODEs was adapted so that the force of infection was dependent on the number of individuals in the home range of the recipient species, rather than the density of hosts in a km^2 .

Results

Null disease model results (R_0 constant across all species). Assuming that host competence does not vary between species, we simulated how changing the size of a habitat patch, thus changing the diversity of hosts, would affect pathogen prevalence. For frequency-dependent transmitted pathogens, $\left(\frac{\beta SI}{N} \right)$, additional species decreased the prevalence because contacts were divided between all hosts (Fig 1, column 2) and the addition of larger bodied species meant that contacts were likely to be with individuals that had a lower force of infection in the community (Fig S5C). By contrast, for pathogens with density-dependent transmission $\left(\frac{\beta SI}{A} \right)$, additional species increased prevalence within each species and across the community, irrespective of the new species' body size relative to the existing community (Fig 1, column 3). The largest changes in equilibrium prevalence of a disease occurred when subsequent species were lost in small and intermediate habitat patches, mainly because these species changed the density and number of individuals in the habitat patch to a greater degree than large bodied species. This was especially true when between species transmission was very small (i.e. 10%) compared to within species transmission (Fig S6). These outcomes assumed that R_0 , and therefore host competence, did not depend on host body size. In these simulations, we assumed that the disease does not cause mortality ($\alpha = 0$).

Changing assumptions regarding the order of community disassembly did not affect the general relationship between diversity and disease - as long as species abundance and presence were nested. Although the general trend did not change for nested community assemblies, the rate at which prevalence within each species and in the overall community increased (DD) or decreased (FD) with habitat size and as host density increased (Fig 1, Row 2).. When the densities were not constant across habitat size, the relationship between prevalence, species richness, and habitat size were less predictable (Fig 1, Row 3). Although on average the relationships between diversity and disease were consistent among pathogen transmission types, there is more variation in prevalence (Fig S7).

When host competence was dependent on body size (R_0 varies among species). If R_0 within populations of smaller species was larger than R_0 within populations of larger-bodied species [54], then the difference in prevalence of a frequency-dependent pathogen between the smallest and largest habitat was even greater than observed for null simulations (Fig S8). This means when smaller hosts are more competent at transmitting frequency dependent pathogens, the dilution effect should be easier to observe. Yet, when the same assumptions are applied to density dependent pathogens (Fig. 2A), there is still an amplification effect. Even though a higher proportion of competent hosts occupy smaller habitats, larger habitats have a higher host density, and thus drive higher pathogen prevalence, at least within the range of R_0 used in these scenarios.

Under the assumption that behavioural traits vary with body size and influence R_0 for a given pathogen [59], biodiversity can increase disease risk for frequency-dependent pathogens at larger more diverse patch sizes, despite a dilution effect between small and intermediate patches (Fig. 2B). The assemblage of species and a significant change in R_0 by body size (Fig S5) can reverse the relationship between biodiversity and prevalence observed for null models.

When we assumed that host competence was unrelated to body size and dead end hosts were randomly assigned to species, there were communities that produced non-monotonic changes in prevalence across habitat loss. For each community simulation, 50% of species in the global community were assigned dead-end host status. These dead-end hosts could become infected but were not infectious. When the smallest host was

incompetent, the pathogen invaded only when competent hosts were present in habitat patches. Incompetent hosts always decreased the prevalence of disease when they are added to a community with frequency-dependent pathogens (Fig. 2C). For frequency dependent pathogens, the distribution of body sizes of the incompetent species determined the degree of this decline, but decreases could be 'rescued' by the addition of competent host species (Fig. S9).

Heterogeneous mixing of populations. If we assumed host contacts scaled with home range size, then the direction of diversity-disease relationships remained unchanged. However, when new host species entered habitat patches, the within-species prevalence increased more than in the baseline assumptions (Fig 3B vs. Fig 3A). The greater infection prevalence in these hosts was driven by a larger home range, and thus more potential infectious contacts over the lifetime of an individual. While individual species had greater infection prevalence, the community infection prevalence was lower than the baseline model (Fig. 3A). As body size increased, transmission rate declined for a density-dependent model including home ranges of species. Prevalence increased because the overall density of hosts was greater in species-rich patches, but the larger bodied species were less efficient at transmitting pathogens and thus the increase in prevalence was less steep than in the baseline simulations.

Discussion

We simulated null expectations of changing disease prevalence in a host community in shrinking habitat. Using an allometric multi-host model, we showed that declining biodiversity can either amplify or dilute pathogen prevalence. The direction of the effect depended upon the transmission mode and how relative host competence scaled with body size. Amplification effects occurred when we increased host diversity for pathogens with density-dependent transmission. Dilution effects were observed for simulations when host diversity increased for pathogens with frequency dependent transmission. In contrast, dilution effects were never observed for density-dependent pathogens when host competence varied among species. Incorporating assumptions of home ranges and community composition did not affect diversity-disease relationships for a given pathogen transmission mode, but did change the magnitude of increases in prevalence observed over the community. This framework we have developed can be tailored to specific systems and host assemblages to predict disease prevalence in habitats of varying size. It can also be used to determine the assumptions necessary for observing specific diversity-disease relationships.

Allometric scaling provides an appropriate tool for understanding null expectations for diversity and pathogen transmission across a fragmenting landscape. Allometric disease models can recreate epidemic cycles observed in nature [60, 61] and empirical evidence supports scaling of disease parameters by body size [62]. Although another study also suggested host competence affects the relationship between diversity and disease [54], we show that these difference are significant (up to a 40% change in prevalence) and show that this can be detected in field situations. Furthermore, the proposed allometric model across declining habitat produced results that are consistent with empirical observations of pathogens in hosts experiencing habitat loss. For example, the prevalence of New World *Trypanosoma* species (vector-borne protozoans) in primate [37] and bat [40] populations in habitat patches is higher than the prevalence in continuous habitats where host diversity and average biomass is higher. The presumed density-dependent fungal pathogen, *Batrachochytrium dendrobatidis*, the cause of chytridiomycosis in amphibians, has a lower prevalence in hosts that reside in small habitat patches with lower amphibian diversity than in hosts that reside in contiguous habitat [63]. The models we presented simulated prevalence across a range of habitat sizes, but the largest differences were between patches where 1-3 host species were present, compared with the full host community (10 host species). In field studies, differences in prevalence are likely to be observed, but significant differences may only be detectable between communities that have a large difference in species richness.

Confirming results of other theoretical work, we found that transmission mode was consistently the best predictor of diversity-disease relationships. We recognize that these transmission processes are an over simplification of actual dynamics [52], but they provide an important benchmark and models with these frameworks can be used to explain empirical data. Density dependent multihost pathogens may best capture dynamics of foot and mouth disease (Tildesley et al 2007). Frequency dependent transmission has better represented dynamics of lyme disease (*Borellia burgdorferi*), sarcoptic mange, and *Mycoplasma ovipneumoniae* [14, 64, 65]. In contrast, work on other wildlife pathogens show that empirical data are best represented by models

that incorporate transmission that lies in between frequency - and density-dependent transmission [66] et al 2009, Cross et al. 2013). If the

The patterns of community assembly and disassembly are important for disentangling diversity-disease relationships. Previous work assembled communities from a global species pool [67-69], using Fisher's Law and Preston's Law for determining the number of vector and host species, respectively. In our models, we are considering a community within shrinking habitat with predictable community disassembly patterns. Although fragmentation can have complicated effects on local species abundance and persistence, overall habitat loss consistently excludes larger bodied species from landscapes. Our model is consistent with empirical observations: larger habitats support a greater number of species than smaller habitats; larger habitats support increasing numbers of larger-bodied species [70, 71], all at higher net density. Nestedness of host communities has been recorded in several systems - suggesting that the hosts found in the least biodiverse communities are predictable [72, 73].

A key research focus should be how to quantify host competence across body size and determine if predictable changes occur across multiple taxonomic groups [73]. If we are also able to quantify how host competence varies across the habitat range, then we will improve predictions of how diseases respond to changes in habitat. Changes in host competence may be even more important than variations in host density [73], which are often highlighted as important factors driving disease dynamics. Additionally, we show that these differences in competence can lead to dilution effects and amplification effects, depending on the details of habitat loss and community assembly.

Host competence can be described as the product of the probability of getting infected (susceptibility), the duration of infectious period, and the probability of infecting another host. Host competence varies in natural systems [74-76], and this variation has been highlighted as an important mechanism driving the dilution effect in natural systems [54, 55, 72, 73, 77, 78]. Our simulations confirm the importance of host competence- although we did not explicitly include each component of host competence; we varied host competence by changing species-specific R_0 , resulting in a change in transmission rate (β_{ii}) but not infectious period ($1/\gamma$). By linking host competence to body size, we demonstrated that the amplification effect could be observed for frequency-dependent pathogens- reversing the effect observed when host competence is assumed to be constant across species. Fast-lived species (i.e. smaller bodied size) are often thought to have higher host competence than larger bodied hosts. For host competence to affect transmission of density-dependent pathogens, additional incompetent hosts must also be competitive and suppress or indirectly compete with the competent hosts. Unless the density of competent hosts changes, the addition of species will not affect dynamics even if they are incompetent. Competition may explain why the dilution effect has been observed in density dependent pathogens such as Sin Nombre virus [10, 79]. Although the dilution effect was observed for frequency dependent pathogens using null assumptions, decreasing host competence with larger body size increased the dilution effect observed. Therefore, incompetent hosts (also called incidental hosts), facilitated a dilution effect (i.e. [80]), but their inclusion was a sufficient, but not necessary, condition to explain the dilution effect for frequency dependent pathogens.

Alternatively, it has also been proposed that R_0 is greater in larger-bodied species. This general trend is driven by increases in average group size as hosts get larger. If larger hosts are actually more competent, this exacerbates the amplification effect observed for null expectations of density-dependent pathogens. It can also flip the null observations of frequency dependent pathogens, so that in extreme cases the amplification effect can be observed. Although we do not know of any empirical examples to support this, the lack of published results may be due to publication biases [23] rather than absence of this phenomena in natural systems. Variation in host competence may arise from vector preference for particular hosts [81], rather than innate host biology [77], which in turn may be related to allometry. Understanding how host species contribute to infection dynamics [82] and how this changes with body size, and thus community assemblage, should be a priority for future empirical studies.

Most of our simulations indicated a monotonic change in prevalence and diversity - either increasing (amplification) or decreasing (dilution) prevalence as patch size and diversity increased. Yet some field studies show a concave relationship between prevalence and diversity, so that the highest observed prevalence is found in intermediate fragments and the lowest prevalence is found in the smallest and largest habitat fragments [37, 41]. These situations may arise when the community abundance peaks prior to biodiversity as

habitats get larger- these conditions were simulated using similar sets of differential equations [33]. These patterns could also in our scenarios when larger hosts are more competent (Fig 2B), mirroring trait-based amplification effects observed for frequency dependent pathogens by O'Regan and colleagues [50]. Competent hosts that enter intermediate habitat patches could bolster infection prevalence and subsequent addition of incompetent hosts in larger habitats could decrease prevalence, but this is complicated by the size distribution of hosts and their relative densities. Host and pathogen life history could also play a role in individual parasite species' responses to habitat fragmentation [43, 44], particularly for pathogens with complex life cycles. These complexities underscore the importance of understanding how species assemble and how host competence changes with increases in diversity.

We did not include explicit models of vector populations, which are expected to undergo frequency-dependent pathogen transmission. Including non-competent hosts into vector-borne disease models reduces disease prevalence as biodiversity increases [24]- yet observed dilution effects in populations that undergo frequency dependent transmission do not necessarily require the addition of incompetent hosts, as it is the null expectation for this type of transmission. Temporal and spatial heterogeneity within a habitat patch [83], host distributions [84], and immunological responses to prior infection [85] will also impact transmission dynamics of vector-borne pathogens. Stage-specific transmission dynamics have also been shown to impact disease dynamics [86]. Incorporating vector populations into these null models and how these respond to changes in habitat area and host availability in these models will be an essential next step.

Although habitat loss often results in a reduction in overall diversity, effects on individual species abundance and distribution are often vary [87]. Patterns of infection in wildlife populations will be influenced by host ranging patterns, density, predation, intraspecific and interspecific contact rates and diet [88], and these characteristics can be affected by changes in habitat structure [89, 90], edge effects [91], and synergistic effects of habitat loss and disturbances such as fire [92]. The presence of competitors and/or predators has been suggested as a mechanism for reducing infection prevalence in communities with high levels of biodiversity [24, 93] and is supported by a model with varying interspecific contact rates[50]. These variations in species-specific responses to habitat loss will be important to consider for certain disease systems and will likely influence changes in prevalence depending on the host competence and sensitivity to habitat loss. Here we highlight that host communities decline in a predictable manner, emphasizing a nested host species community that is dominated by smaller bodied species in low diverse, small habitats. While this produces the dilution effect for frequency dependent pathogens and the amplification effect for density dependent pathogens, changes in how host competence scales with body size can alter these expectations.

Our multi-host model focuses on a single habitat patch and describes infection dynamics that vary with patch size and community composition. In natural populations, habitat patches are rarely isolated to the extent that immigration and emigration are non-existent. Habitat patches exist in meta-populations with varying levels of connectivity and models emphasize that connectivity and host dispersal are important characteristics determining system dynamics [5, 94-96]. Larger habitat patches may support higher incidences of density-dependent pathogens and serve as sources of pathogen pollution for small habitat fragments in the landscape. In contrast, if movement only occurs towards larger, higher quality habitat, the average prevalence in large patches could decline if uninfected individuals emigrate, or increase if the density of susceptible hosts increases. The framework we present here could be extended to incorporate spatial structure so that the influence of allometric home ranges and dispersal on prevalence can be explored in each sub-population within a meta-population/meta-community context [59].

Critically, the temporal and spatial scale at which observations are made will affect the infectious disease dynamics [26]. Moving forward, some of the biggest challenges will be to determine at which spatial and temporal scale empirical measurements would be useful for testing these null predictions. Disease prevalence should be measured in fragmented long-term ecological study sites, such as Stability of Altered Forest Ecosystems (S.A.F.E.) in Borneo [97] or Biological Dynamics of Forest Fragments Project (B.D.F.F.P.) in Brazil [98]. These systems have existing data on host demographics, population densities, community compositions, and environmental changes, which would complement disease dynamic investigations. Changes in pathogen prevalence can be observed with respect to transmission mode, pathogen characteristics, and host competence in order to disentangle mechanisms driving transmission dynamics.

Conclusion

Using realistic allometric models of a multi-host pathogen, we showed that amplification and dilution effects can be observed in a shrinking habitat. The observed change in equilibrium prevalence with declining habitat and decreasing biodiversity depends on how communities disassemble, how competence scales with body size, the likelihood of a given species' residence in a patch, and the transmission mode. The infectious disease-habitat loss model developed in this study provides a useful template for the design of longitudinal empirical studies of multi-host pathogens in shrinking habitat. It can be adapted to study-specific host species, transmission mode, and habitat loss responses to direct sampling effort across a landscape.

Understanding how habitat loss, biodiversity, and disease are related is an essential challenge in natural area management. In cases where habitat loss correlates with emerging diseases of humans [99-101], management of disease systems will benefit from evaluating whether biodiversity is the underlying mechanism of pathogen emergence, or if it is a combination of changing contact patterns, environmental conditions, and complex species interactions driving emergence. While we often focus on human health, habitat loss and subsequent changes in disease incidence also affect disease risk in animal and plant populations [5, 102].

Additional Information

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Ethics

No ethical approval was required.

Data Accessibility

The data gathered model parameterization was gathered from the primary literature. All data has been included as tables or figures in the Supplementary Information.

Authors' Contributions

LSPB APD CLF TRG NLG HIM and RKP conceived the study. CLF APD RKP and NLG designed the study and analyzed the results. CLF NLG LSPB and TRG acquired empirical data for model conceptualization. CLF RKP NLG APD LSPB MDW HIM and TRG wrote the manuscript and all contributed to the editing and approval of the final draft.

Competing Interests

We have no competing interests.

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Figure captions

Figure 1. Community disease prevalence in a shrinking habitat. The prevalence of infection in individual species and the overall community at equilibrium ($t=150$ years) in both frequency-dependent (column 2) and density-dependent (column 3) transmission simulations are shown. There was no variation in host competence or between species contact (within species $R_0 = 2.0$; $\psi = 0.5$). Simulations for a single community, shown in color, represent a community with species that have an average mass of 0.011, 0.030, 0.065, 0.075, 0.23, 0.537, 1.505, 1.515, 13.333, 14.201 kg. For all simulations, the prevalence for each species is shown with colors representing their size, from brown (smallest host) to aquamarine (largest host). The black line indicates the prevalence of disease across the community and is most similar to the intermediate body classes.

Simulations with 100 random communities are shown in Fig. S7. Each row in the figure represents a unique community assemblage assumption (detailed in methods). A pathogen with frequency dependent transmission declines in prevalence as habitat area increases (from left to right) and species richness increases, thus leading to a dilution effect, although the strength of this declines with increasing randomness in the community structure. Pathogen prevalence within each species and across the entire population increased as habitat size increased for density dependent pathogens, demonstrating an amplification effect, however this asymptotes when species were no longer added to the community. Community assembly only affects the rate of increase or decrease in prevalence when species and abundance are nested, but not the directionality of diversity-disease relationships. When species presence and density are not directly related to area, then the relationship between diversity and disease becomes less predictable.

Figure 2. Variation in host competence and underlying assumptions impact diversity- disease relationships. **A.** Even if the smallest hosts are the most competent [73] and there are extreme differences in R_0 between body sizes (Fig S5), only the amplification effect is observed for density dependent pathogens. **B.** If behavioural allometry leads to an increase in R_0 across body size, this can lead to an amplification effect for frequency dependent pathogens at larger patch sizes, but a dilution effect for small to intermediate patches. **C.** When species that can become infected but are in turn not infectious (incompetent hosts, denoted by x) are randomly assembled along the distribution of body sizes, then dilution effects can be exacerbated for frequency dependent pathogens, but this depends on the order of community introduction of these incompetent hosts.

Figure 3. Impact of heterogeneous contacts on density dependent pathogen transmission. Using identical disease parameters, endemic prevalence was observed for a host system that assumed contacts were determined by density of hosts (A) and compared to a system in which home range determined the average contacts of an individual from a given species (B). When home range is not taken into account, overall prevalence is higher, but when home range is considered larger bodied species that have larger home ranges have higher within – species infection prevalence.