



Species diversity concurrently dilutes and amplifies transmission in a zoonotic host–pathogen system through competing mechanisms

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In this era of unprecedented biodiversity loss and increased zoonotic disease emergence, it is imperative to understand the effects of biodiversity on zoonotic pathogen dynamics in wildlife. Whether increasing biodiversity should lead to a decrease or increase in infection prevalence, termed the dilution and amplification effects, respectively, has been hotly debated in disease ecology. Sin Nombre hantavirus, which has an ~35% mortality rate when it spills over into humans, occurs at a lower prevalence in the reservoir host, the North American deer mouse, in areas with higher small mammal diversity—a dilution effect. However, the mechanism driving this relationship is not understood. Using a mechanistic mathematical model of infection dynamics and a unique long-term, high-resolution, multisite dataset, it appears that the observed dilution effect is a result of increasing small-mammal diversity leading to decreased deer mouse population density and, subsequently, prevalence (a result of density-dependent transmission). However, once density is taken into account, there is an increase in the transmission rate at sites with higher diversity—a component amplification effect. Therefore, dilution and amplification are occurring at the same time in the same host–pathogen system; there is a component amplification effect (increase in transmission rate), but overall a net dilution because the effect of diversity on reservoir host population density is stronger. These results suggest we should focus on how biodiversity affects individual mechanisms that drive prevalence and their relative strengths if we want to make generalizable predictions across host–pathogen systems.

dilution effect | amplification effect | hantavirus | SIR modeling | zoonotic disease

Biodiversity is being lost at an unprecedented rate; some say we have entered the sixth mass extinction (1). Meanwhile, zoonotic infectious disease outbreaks are increasing (2, 3). Therefore, it is imperative to understand how biodiversity affects disease dynamics in wildlife. Work on Lyme disease demonstrated how, under certain conditions, increased vertebrate species diversity can lead to decreased disease transmission and prevalence in the reservoir host and subsequently decreased spillover to humans—the so-called “dilution effect” (4, 5). The generality of the dilution effect has been the topic of much recent debate. If it is a general phenomenon, preserving biodiversity would be a win–win for animal conservation and control of zoonotic diseases. Mounting evidence suggests the dilution effect applies to several systems across taxa (6–8). However, the dilution effect does not appear to be universal but depends on the animal community composition, host and pathogen ecologies, and the scale at which the system is examined (9–12). In some circumstances, increased species diversity can lead to increased infection prevalence, the “amplification effect” (10–13). For most systems, exactly which circumstances and mechanisms lead to disease amplification or dilution have

not been fully characterized, particularly for directly transmitted diseases (14).

Much of the research on this issue has focused on vector-borne (e.g., tick or mosquito-transmitted) or environmentally transmitted diseases, such as Lyme disease, West Nile virus, and *Ribeiroia ondatrae*, where the dilution effect is caused by more diverse communities having multiple species with a range of competencies with respect to pathogen replication and vector preferences (5, 15, 16). In these systems, when biodiversity is lost, the species that remain tend to be the most competent reservoirs. [The generality of this phenomenon is debated (13, 17).] Directly transmitted diseases have received little attention.

For directly transmitted pathogens with one main reservoir host, the SIR (susceptible–infected–recovered) framework is useful for conceptualizing how species diversity could affect infection dynamics. These models are called compartmental models, because individuals within a population are grouped into different compartments with respect to their disease status. Individuals are either uninfected and susceptible to infection (*S*), infected and infectious (*I*), or recovered and immune (*R*). Over time, individuals flow through these compartments. The rate of new infections is determined by the density of susceptible individuals (*S*), the density of infected individuals (*I*), the transmission rate (β), and whether contact rates increase with density (density-dependent transmission) or are independent of

Significance

There has been an impassioned debate in recent years about whether biodiversity is negatively or positively correlated to wildlife and zoonotic disease transmission and risk, suggesting a “dilution” or “amplification” effect, respectively. Here, we demonstrate for an important zoonotic disease (hantavirus pulmonary syndrome) that species diversity can act differently on competing drivers of disease transmission (host density, contact rates, transmissibility) and may cause increases and decreases in transmission, concurrently in the same host–pathogen system. The net effect (dilution, amplification, or no effect) is determined by the strength of the competing mechanisms. Therefore, to move forward, researchers should focus on how biodiversity affects individual mechanisms separately and their net effects if we want to make generalizable predictions across systems.

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density (frequency-dependent transmission). Therefore, the rate of new infections is βSI or $\beta SI/N$, for density-dependent or frequency-dependent transmission, respectively.

The transmission rate, β , is a product of the contact rate and the probability of transmission given contact (or transmissibility). Therefore, three mechanisms determine how diversity could affect pathogen transmission and prevalence. Dilution will occur if increased species diversity leads to (i) decreased host population density (if transmission is density-dependent), (ii) decreased contact rates, and/or (iii) decreased transmissibility (Fig. 1, blue). Conversely, amplification could occur if increased diversity leads to increased host density, contact rates, and/or transmissibility (Fig. 1, red).

Sin Nombre hantavirus (SNV) is a directly transmitted zoonotic pathogen whose reservoir host is the North American deer mouse (*Peromyscus maniculatus*, hereafter deer mouse). Humans are spillover hosts infected when they have direct contact with urine, feces, or saliva from an infected host rodent or inhale infectious aerosols of these substances. In humans, SNV causes hantavirus pulmonary syndrome (HPS), which has an overall 36% mortality rate. SNV prevalence in deer mice is often lower in communities with higher small mammal diversity—the dilution effect (18–20).

Because SNV is a directly transmitted disease (no vector) that is highly host-specific and contacts of infected deer mice with members of other species in the community are not likely to pass on the infection, all other hosts are essentially noncompetent. Spillover of SNV from deer mice may occur (e.g., to other rodents and humans). However, they are not likely to be important in the reservoir dynamics except under unusual conditions (21). Thus, the mechanism for how small mammal diversity acts on SNV prevalence must be different from Lyme disease, West Nile virus, and *R. ondatrae* and has yet to be determined. The SIR framework with one reservoir host is appropriate for examining this question. Accordingly, the decrease in prevalence as small mammal diversity increases must be caused by at least one of the following mechanisms: (i) decreased host population density (N), or (ii) decreased transmission rate (β) within the host population by either decreasing (a) intraspecific contact rates and/or (b) transmissibility of the pathogen.

For the first mechanism, host population density, to be the driving factor, there must be density-dependent transmission, which leads to decreased infection prevalence with decreased deer mouse density. Some researchers concluded host density could not be the mechanism driving dilution because they found no simultaneous positive relationship between deer mouse population density and SNV infection prevalence (19, 20). However, we recently clarified the density–prevalence relationship showing that in host–pathogen systems with highly variable dynamics (where the carrying capacity and host density vary widely over

space and time), the system never reaches equilibrium. Therefore, host population density can have a strong effect on SNV prevalence but with time lags that themselves depend on density, thus making the relationship not apparent using statistical correlations (22). Due to these nonequilibrium dynamics, the relationship between density and prevalence can only be deciphered using a dynamical mathematical model of disease dynamics. We showed SNV does have density-dependent transmission (22); therefore, reduced host density could, indeed, be driving the dilution effect.

Alternatively, the dilution effect could be caused by increased small mammal diversity decreasing the transmission rate (β), by decreasing the contact rate or the transmissibility of the virus between mice.

To determine the mechanism of the observed dilution effect for SNV, we examine the two components that determine prevalence (density and transmission rate) by fitting a mechanistic model to data from a multistate long-term study of hantavirus ecology. The US Centers for Disease Control and Prevention (CDC) and other sources funded longitudinal monitoring of small mammal communities at more than 40 sites in Arizona, Colorado, Montana, and New Mexico. Starting in 1994, rodents were trapped monthly at all sites (except every 6 wk in Colorado) for up to 15 y. To synthesize drivers of infection across sites with differing dynamics, we apply the same dynamical modeling framework to examine all sites with a time series of infection long enough to analyze. We apply a mechanistic dynamical SIR-type model that simulates infection dynamics given host density (22) to 18 sites in 4 US states, with varying species diversity, estimating a transmission rate for each site.

We hypothesize that the dilution effect is driven by host population density—sites with higher small mammal diversity have lower deer mouse density, and SNV prevalence is lower as a result of density-dependent transmission. This leads to the prediction that when we fit our mechanistic model, driven by host density, to each site, the model will fit the observed host–pathogen dynamics, and the transmission rate (β) will not vary as a function of diversity. The alternative hypothesis is that the observed dilution effect is driven by contact rates and/or probability of transmission given contact, components of the transmission rate (β). This predicts that when we fit our model to multiple sites, the estimated transmission rate (β) will be lower for sites with higher species diversity.

Materials and Methods

Study Sites. We analyze data from small mammal capture–mark–recapture SNV studies in Montana, New Mexico, Arizona, and Colorado (Fig. 2). Data span June 1994 through 2015, with sites trapped an average of 79 mo (range 3 to 179). *SI Appendix, Table S1* summarizes data from all 40 sites. We applied our model to sites in which there was an average of >2 deer mice/hectare, >4% SNV prevalence, and >12 mo of data (18 sites, Table 1); because our model is dynamical and prevalence depends on previous population density, we excluded sites with too few data points.

In Montana, 10-by-10 trap grids contained 100 Sherman ($8 \times 9 \times 23$ cm; H.B. Sherman Trap Company) live-capture traps at 10-m intervals covering 1 ha. See ref. 23 for details. Grids were trapped monthly as weather allowed. In the SW (Arizona, Colorado, New Mexico), trapping webs were used rather than grids. Each web covered 3.14 ha and contained 12 100-m transects radiating from a central point. Each web contained 148 Sherman traps. Webs in Arizona and New Mexico were trapped monthly and in Colorado every 6 wk, as weather allowed. See *SI Appendix, Table S1* for details about frequency and duration of trapping at sites, deer mouse population densities, SNV prevalence, and small mammal diversity. See ref. 24 for more details of sites and protocols.

Trapping protocols were similar across sites. As a rule, researchers trapped for three consecutive nights. Traps were set in the evenings baited with peanut butter and oats, cracked corn, or mixed grain. Cotton or polyester fiberfill was added for nest material. Traps were checked in the mornings. Captured small mammals were anesthetized, ear-tagged, weighed, and a

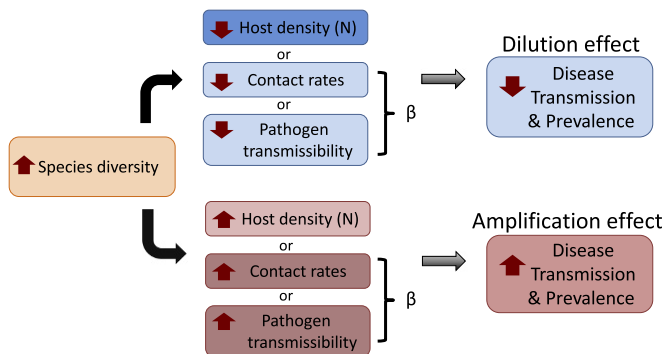


Fig. 1. Conceptual diagram of how species diversity could affect transmission and prevalence of a directly transmitted disease.

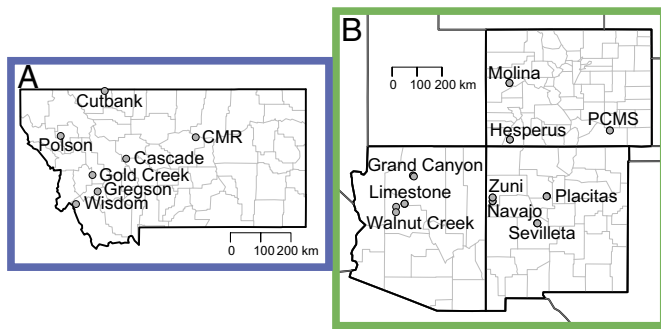


Fig. 2. Map of study sites in (A) Montana and (B) the southwest (SW) United States, including Arizona, Colorado, and New Mexico.

blood sample taken to test for SNV antibodies. Because SNV is a chronic infection in deermice, antibodies are an appropriate indicator of current infection. Serologic testing was conducted at CDC, Atlanta; the Montana Department of Public Health and Human Services; Montana State University; or Montana Tech using a standard enzyme-linked immunosorbent assay (ELISA) for IgG antibody to SNV (25).

We analyze each web/grid separately because often they were trapped for differing lengths and were sufficiently distant from one another that many of the grids' community dynamics were not significantly correlated. Additionally, we group Montana and SW sites separately because of the different trapping structure and minor differences in data collection. For comparing data across Montana and the SW, we normalize density to 1 ha.

Model. Deermouse population dynamics are highly variable over time and strongly tied to local environmental conditions (26, 27). We have developed a mechanistic mathematical model of SNV in deermice that allows for the environmental carrying capacity and population density to vary over time (essential in this highly variable, bottom-up system) (22). Here we use a version of our model, which is a type of SIR compartmental model; SNV is a chronic, probably life-long infection in deermice, and thus, there is no *R* class. The model includes logistic growth, where the host population experiences density dependence partitioned across birth and death rates and determined by a time-varying carrying capacity, K_t . The underlying model is

$$\begin{aligned} \frac{dS}{dt} &= N \left(b - ar \frac{N}{K_t} \right) - S \left(d + (1-a)r \frac{N}{K_t} \right) - \beta SI & [1] \\ \frac{dI}{dt} &= \beta SI - I \left(\mu + d + (1-a)r \frac{N}{K_t} \right) + \phi, \end{aligned}$$

where *S* is the density of susceptible individuals, *I* is the density of infected individuals, and $N = S + I$. [This is our published model (22), except without age structure, which did not significantly impact dynamics.] *b* is the maximum birth rate (in absence of density dependence; i.e., when $N = 0$), *d* is the minimum death rate, and *a* is the proportion of density dependence due to density dependence in birth rates. If $a = 0$, birth rates are density independent, and all of the density dependence is on the death rates; if $a = 1$, all of the density dependence is on the birth rates. *r* is $b - d$. μ is the disease-induced mortality rate. β is the transmission rate, and ϕ is the immigration rate of infected individuals (see *SI Appendix*, Fig. S1 for life-cycle diagram).

This is the underlying model we use; however, for the purposes of our analyses—to understand the importance of deermouse population density in driving dynamics—we are interested in predicting *I* assuming we know the population density, *N*. Therefore, we only need to model *I*, since $S = N - I$.

Previously, we estimated parameters for study sites in Montana (22, 28). Here we assume that the disease-induced mortality rate (μ), maximum birth rate (*b*), minimum death rate (*d*), and how density dependence is partitioned (*a*) are the same across all sites [$b = 0.315$, $d = 3.66 \times 10^{-5}$, $a = 0.614$, and $\mu = 0.085$ (22)]. However, the environmental carrying capacity (K_t) and therefore the realized birth and death rates vary across sites and over time at the same site. The quantities that vary across sites include the inputs for the model: *N*, K_t , and initial infected deermouse density (I_0), all taken from the data. Parameters, transmission rate (β) and infected immigration rate (ϕ), are estimated using maximum likelihood for each site.

Parameter Estimation. We previously found that setting K_t to a smoothed spline of the deermouse population density 3 mo ahead approximated the time-varying carrying capacity (density lags behind carrying capacity by 3 mo) (22). Using this approximation, the model-predicted deermouse population dynamics matched observed deermouse population density. Needing future deermouse density to predict current infection levels makes this implementation of the model less useful for prediction. However, our goal is to understand what processes are leading to the observed dynamics, not prediction.

As an index of relative population density, we use minimum number known alive (MNA) (29). Similarly for relative density of infected individuals, we use minimum number known infected (MNI). See *SI Appendix* for justification of this index and comparisons with robust design population estimates that account for probability of detection using Program Mark.

We estimate a transmission rate (β) and infected immigration rate (ϕ) for each site separately using trajectory matching (30). For this, we numerically minimize the negative log likelihood between the vector of model-predicted density of infected mice and the vector of observed MNI using the Nelder–Mead algorithm implemented in the “optim” function in R, assuming Poisson errors. The only inputs are initial *I* (set to that observed in the data) and N_t and K_t (both smoothed splines from the data to accommodate the continuous-time framework).

We then compare the estimated transmission rate (β) for each site to the average small mammal diversity for that site, using Simpson's diversity index, $D = 1 / \sum_{i=1}^R p_i^2$, where p_i is the proportional abundance of species, *i*, and *R* is the small mammal species richness—that is, total number of small mammal species trapped at that site. We also compare results to the Shannon–Weiner index and species richness (see *SI Appendix*).

See *SI Appendix* for data and R code.

Results

We observe a dilution effect—a decrease in average SNV prevalence in deermice with increased average small mammal diversity—in the SW ($P = 0.002$, $R^2 = 0.64$; Fig. 3A) but not in Montana ($P = 0.60$; Fig. 3B). However, our dynamical epidemiological (*SI Appendix*) model, which simulates infection based on deermouse density, fits both Montana and SW sites well (Fig. 4 and *SI Appendix*, Figs. S2 and S3). Once density is taken into account using the model, the estimated transmission rate (β) is significantly correlated to diversity. However, it is in the opposite direction to what is predicted if the dilution effect is driven

Table 1. Summary of longitudinal data from modeled sites

State	Web or grid	Mo.	Dens.	Prev.	<i>D</i>	β
MT	Cascade.11	179	25.65	0.048	1.06	0.0040
MT	CMR.18	69	8.70	0.044	1.51	0.0183
MT	Cutbank.15	70	10.59	0.152	1.52	0.0202
MT	GoldCreek.8	70	11.24	0.133	1.62	0.0091
MT	GoldCreek.9	70	8.60	0.086	1.86	0.0152
MT	Gregson.Upper	134	10.49	0.137	1.01	0.0106
MT	Gregson.Lower	158	16.18	0.108	1.06	0.0073
MT	Polson.5	96	53.64	0.213	1.12	0.0027
MT	Polson.6	70	14.91	0.125	1.54	0.0084
AZ	Grand Canyon.E	43	20.15	0.277	1.72	0.0042
AZ	Grand Canyon.M	46	9.57	0.311	1.52	0.0120
AZ	Grand Canyon.T	46	8.73	0.152	1.80	0.0184
CO	Hesperus.ha	67	8.43	0.172	1.77	0.0138
CO	Hesperus.hb	68	9.28	0.183	1.34	0.0141
CO	Molina.ma	58	4.84	0.062	2.53	0.0277
CO	Molina.mtb	47	4.49	0.099	2.21	0.0326
NM	Navajo.1	92	3.27	0.065	2.47	0.0523
NM	Zuni.2	92	2.06	0.089	3.76	0.0789

AZ, Arizona; β , estimated transmission rate, per hectare; CMR, Charles M. Russell Wildlife Refuge; CO, Colorado; *D*, Simpson's *D* diversity index; dens., average deermouse density, per hectare; mo., number of months sampled; MT, Montana; NM, New Mexico; prev., average SNV antibody prevalence. See *SI Appendix*, Table S1 for more information and all sites.

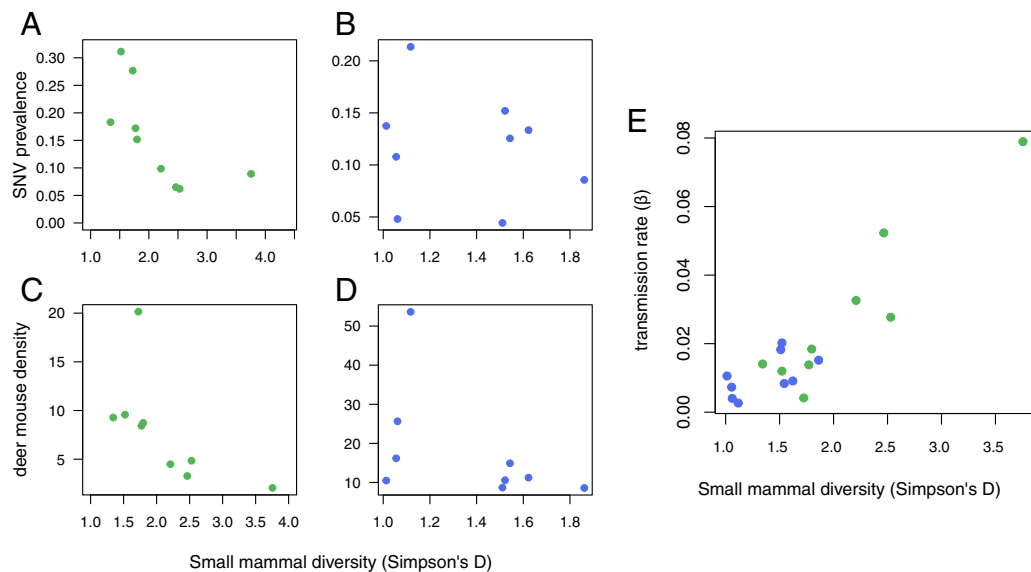


Fig. 3. Relationship between average small mammal diversity by Simpson's diversity index and (A) average SNV prevalence in deermice for SW sites and (B) Montana sites, (C) average deermouse population density for SW sites and (D) Montana sites, and (E) estimated transmission rate (β) among deermice. Throughout, SW sites are green, and Montana sites are blue.

by host contact rates or transmissibility. There is a positive relationship between diversity and transmission rate—a component amplification effect (Fig. 3E).

An increase in small mammal diversity is associated with a decrease in deermouse density in the SW (Fig. 3C) and decrease in transmission and prevalence—the dilution effect (Fig. 3A; as described in Fig. 1, dark blue pathway). However, in Montana, increased small mammal diversity does not lead to decreased deermouse density (Fig. 3D), and there is not a dilution effect (Fig. 3B). However, overall small mammal diversity in Montana is lower than in the SW (see *x* axis for Fig. 3E). Although there is not a significant dilution effect in Montana (because the relationship between diversity and deermouse density is not the same as in the SW), the relationship between transmission rate

(β) and diversity is similar across SW and Montana sites (Fig. 3E). Results are robust to different diversity measures (see *SI Appendix*).

Discussion

This study brings together dynamical ecological models with high-resolution, long-term data, spanning more than 20 y of data collection at 18 sites with different small mammal community composition and deermouse host population dynamics. Our model accurately predicts infection dynamics over time at these disparate sites (Fig. 4 and *SI Appendix*, Fig. S2 and S3), given only deermouse population density and the initial number of infected individuals, and estimating a transmission rate (β) and infected immigration rate (ϕ) for each site. With these data

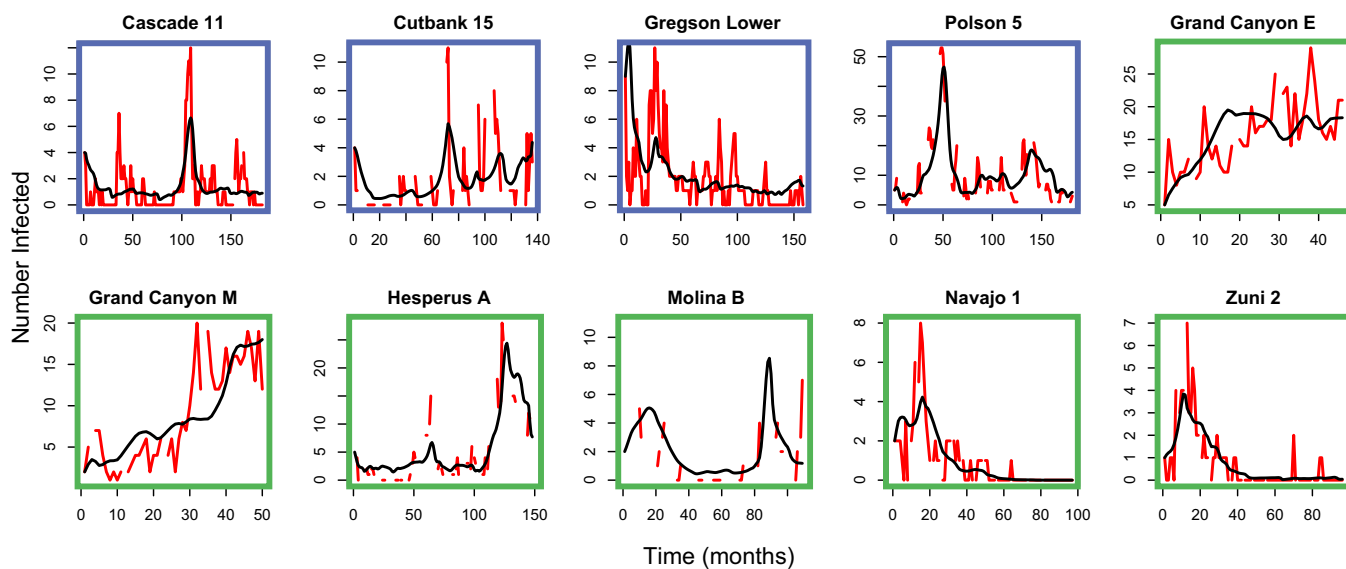


Fig. 4. Fit of models to data from 10 of the 18 sites modeled. Observed number of infected individuals is in red. Model predicted number of infected individuals is in black. Montana sites (blue) are per hectare, and SW sites (green) are per 3.14 ha. See *SI Appendix*, Figs. S1 and S2 for all sites including total deermouse density per site.

and models, we can explore hypotheses about how small mammal diversity affects SNV transmission and disease risk. At our SW study sites, we observe a dilution effect—sites with higher small mammal diversity had decreased prevalence of infection (Fig. 3A). We do not observe a dilution effect at our Montana sites (Fig. 3B).

Given the SIR framework, the dilution effect in the SW must be driven by increased species diversity leading to (i) decreased host population density or (ii) a decrease in the transmission rate, which includes contact rates between deer mice and transmissibility (Fig. 1, blue). If population density (mechanism 1) drove the dilution effect, our model, which predicts infection given host density, should predict the dynamics well, and the transmission rate (β) should not be negatively correlated to diversity. If transmission rate (mechanism 2) drove the dilution effect, our model should predict the dynamics, and each site's estimated transmission rate should be negatively correlated to its diversity. We found support for mechanism 1—that deer mouse population density drives the dilution effect in the SW. Our model fits the dynamics at disparate sites (Fig. 4 and *SI Appendix*, Figs. S1 and S2), and the transmission rate is not negatively correlated to diversity. In fact, it is positively correlated to small mammal species diversity ($R^2 = 0.83$, $P < 0.001$; Fig. 3E). This implies that density causes the observed dilution effect (dark blue, Fig. 1) but that there is an additional component amplification effect through changes in transmission rate (dark red, Fig. 1). Therefore, aspects of both dilution and amplification are occurring at the same time in the same system.

For models with density-dependent transmission, the transmission rate (β) gives the rate of transition from susceptible to infected for a given host density; as density increases, overall transmission increases. Therefore, our finding of increased transmission rates with increased diversity means that for a given host density, transmission occurs at a faster rate in more diverse communities. However, density is a stronger driver of infection dynamics in this system, and since increased diversity leads to lower host density in the SW, there is a net dilution effect in the SW (Fig. 3). In Montana, increased diversity does not lead to a decrease in host density, and there is not a significant dilution effect (Fig. 3).

The dilution effect has been demonstrated for hantaviruses, including SNV in Portland, OR (19) and Utah (20, 31); Puumala virus in bank voles in Finland (32); and hantaviruses (Choclo and Calabazo viruses) in their hosts in Panama (33). However, the mechanism driving the dilution effect has not been fully elucidated for any of these systems. The studies in Finland and Panama reported that increased diversity led to decreased host density and prevalence (32, 33), suggesting density-dependent transmission and host density as a potential mechanism for the observed dilution effect. Neither transmission rate nor its components, contact rates, or transmissibility were measured in these studies. However, the authors of the Portland and Utah studies discounted density as the mechanism for their observed dilution effect, because they did not see a positive (concurrent) relationship between deer mouse density and SNV prevalence and concluded that, therefore, transmission was not density-dependent. However, both studies noted a significant negative relationship between small mammal diversity and deer mouse density (19, 20, 31, 34). Clay et al. (34) found a negative association between intraspecific contact rate and species diversity; however, the relationship may have been largely driven by one outlier. A recent experimental study using enclosures and controlling for density showed no change in intraspecific deer mouse contacts as small mammal diversity increased (35).

For deer mouse population density to drive the dilution effect for SNV, there would need to be density-dependent transmission, which should lead to an observed positive relationship between deer mouse density and SNV prevalence. Field studies

have found mixed results, only rarely showing a concurrent positive relationship (36), and often showing either no relationship (21, 34, 36, 37) or a negative relationship (23, 38). Using our dynamical epidemiological model, we recently clarified how a concurrent negative relationship between deer mouse population density and SNV prevalence can be observed with (positive) density-dependent transmission (22). This negative correlation observed at the Cascade, MT site is a result of the nonequilibrium dynamics (from a constantly changing carrying capacity) in deer mouse population density leading to delayed density dependence, such that prevalence lags density by approximately 8 to 16 mo (22, 39–41). At that site, deer mouse density peaked, and prevalence peaked later when population density had significantly declined again. Note that with delayed density dependence, a positive relationship or no relationship may also be observed between concurrent density and prevalence (e.g., refs. 21, 36, and 37), depending upon the pattern of population fluctuation. We also showed that the delays may not be fixed [e.g., at 1 y (39)] but depend on host density, potentially making the relationship not apparent using conventional statistical analyses or examining averages; it can only be deciphered using a mechanistic dynamical model of infection, such as ours (22). Therefore, we caution against inferring too much from point estimates or averages (e.g., Fig. 3); the true test is how the data match predictions from the dynamical model, which is driven by population density, because it can account for the transient dynamics and inherent variable time lags.

We see a component amplification effect through an increase in the transmission rate at sites with higher small mammal diversity (Fig. 3E). This could be due to increased diversity leading to an increase in contact rates or an increase in the probability of transmission given contact (transmissibility; Fig. 1, dark red). Increased species diversity could increase contact rates between deer mice if, in more diverse areas, deer mice cluster in refugia away from dominant competitors. Alternatively, increased species diversity could lead to increased transmissibility if host animals are stressed in the presence of competitors; increased stress can depress the immune system and lead to an increase in viral replication and increased susceptibility (42). Studies of how contact rates vary with species diversity give mixed results—one study suggested a decrease in contacts (34), but an experimental study (with enclosures controlling for density) showed no change in contacts with increased diversity (35). How transmissibility or immunity is affected by species diversity has yet to be examined and is a plausible mechanism for our observed increase in transmission rates.

Future studies should examine the importance of specific species or what aspects of “biodiversity” lead to both the dilution and component amplification. Host interactions with different species vary. In an enclosure experiment, deer mice had more interspecific interactions with Merriam's kangaroo rats than desert pocket mice or Chihuahuan grasshopper mice, although intraspecific contact rates among deer mice remained unchanged (35). Clay et al. suggested that Ord's Kangaroo rats and pinyon mice may be particularly important in determining SNV prevalence (31).

Although previous studies have shown that the dilution effect can be scale-dependent (11), we show that dilution and amplification can occur at the same time in the same system, at the same scale, and the resultant overall effect will be determined by which effect is stronger. A recent, impassioned debate has centered on whether loss of biodiversity should be more likely to lead to an increase (dilution), decrease (amplification), or idiosyncratic change in disease risk (6–8, 10, 11, 43, 44). Perhaps, the reason that responses to biodiversity seem “idiosyncratic” (10) is because there are differential effects of biodiversity on the different mechanisms driving transmission. Therefore, if we want to make generalizable predictions across systems, we should

focus on how biodiversity affects individual mechanisms that drive transmission and prevalence separately [as called for by, e.g., Johnson et al. (45)] and the relative strength of these.

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