Ecological Responses to Habitat Fragmentation per se

Lenore Fahrig

Geomatics and Landscape Ecology Research Laboratory, Department of Biology, Carleton University, Ottawa, ON Canada K1S 5B6

email: lenore.fahrig@carleton.ca

Abstract

I reviewed empirical studies finding significant ecological responses to habitat fragmentation per se, i.e. significant responses to fragmentation independent of the effects of habitat amount (hereafter, 'habitat fragmentation'). I asked, (1) are most significant responses to habitat fragmentation negative or positive?, and (2) are there particular attributes of species or landscapes that lead to a predominance of negative or positive significant responses? I found 118 studies reporting 381 significant responses to habitat fragmentation independent of habitat amount. Seventy-six percent of these responses were positive. Most significant fragmentation effects were positive, irrespective of how the authors controlled for habitat amount, the measure of fragmentation, the taxonomic group, the type of response variable, or the degree of specialization or conservation status of the species or species group. There was also no support for predictions that most significant responses to fragmentation should be negative in the tropics, for species with larger movement ranges, or when habitat amount is low; most significant fragmentation effects were positive in all these cases. Thus, while 24% of significant responses to habitat fragmentation were negative, I found no conditions in which most responses were negative. Authors suggest a wide range of possible explanations for significant positive responses to habitat fragmentation: increased functional connectivity, habitat diversity, positive edge effects, stability of predator-prey/host-parasitoid systems, reduced competition, spreading of risk, and landscape complementation. A consistent preponderance of positive significant responses to fragmentation implies there is no justification for assigning lower conservation value to a small patch than to an equivalent area within a large patch - just the opposite. It also suggests land-sharing will usually provide higher ecological value than land-sparing.

Key Words: landscape pattern, landscape structure, landscape configuration, landscape complementation, landscape connectivity, landscape heterogeneity, patch area, patch isolation, edge effect, SLOSS

1. HABITAT PATCHINESS

'Habitat patchiness' has been an important concept in ecology for over 80 years, beginning with the observation by Gause (1934) that persistence of predator-prey systems depends on the availability of separate refuges for prey. Huffaker's (1958) classic predator-prey experiment took this idea another step, showing that division of a food resource into a large number of patches allows a predator-prey system to persist, by providing temporary prey refuges that move about in space and time, allowing the prey to stay one step ahead of the predators. Levins & Culver (1971) showed that habitat patchiness can also allow persistence of competing species through a similar dynamic, as long as the better competitor is the worse disperser. And Den Boer (1968) showed more generally that habitat patchiness can increase species persistence by 'spreading the risk' of local extinctions. Thus, from 1934 until about 1970, habitat patchiness was associated with the concept of habitat spatial heterogeneity and was generally considered to have a positive influence on population and community-level ecological responses.

This all changed in 1970 with Levins' extrapolation of the Theory of Island Biogeography (MacArthur & Wilson 1967) to patches of habitat (Levins 1970). From that point to the present, habitat patchiness became associated with the concept of habitat fragmentation and is generally considered to have a negative influence on population and community-level responses (Villard & Metzger 2014; Hanski 2015).

While these two major conceptualizations of habitat patchiness seem to make contradictory predictions, the contradiction is not real. In the earlier work, increasing habitat patchiness (or habitat spatial heterogeneity) generally implied an increase in the total amount of habitat available, or at least no decrease. In contrast, in the later work, increasing habitat patchiness (or habitat fragmentation) generally implied a decrease in the total amount of habitat (Figure 1). Therefore, in the former case the positive effect of patchiness is often associated with habitat gain, while in the latter case the negative effect of patchiness is associated with habitat loss. Thus, both views of habitat patchiness can represent a positive effect of increasing habitat amount on population and community responses.

This then begs the question, what is the influence of patchiness itself on ecological responses, independent of its association with habitat amount? Some have argued that this question is a meaningless academic exercise; habitat loss and fragmentation are so tightly linked that any effort to disentangle their effects is pointless (Didham et al. 2012). However, many examples of environmental policies assume large, negative effects of patchiness independent of habitat amount, i.e., negative effects of 'habitat fragmentation *per se'* (*sensu* Haila & Hanski 1984; Figure 2). For example, in Ontario, Canada, forestry policy specifies that large, contiguous clearcuts can be justified if they "defragment a previous group of smaller cuts" (OMNR 2002). This policy is built on the assumption that large, contiguous blocks of forest have higher ecological value than the *same total area* of small, separated patches of forest (King et al. 1996; Hagan et al. 1997). In general, the assumption that fragmentation has large negative effects on biodiversity implies that policies should emphasize 'land sparing' over 'land sharing' (Green et al. 2005; Figure 2). Are such decisions grounded in fact?

2. STUDY DESIGNS FOR MEASURING EFFECTS OF HABITAT FRAGMENTATION INDEPENDENT OF HABITAT AMOUNT

While habitat fragmentation is often thought of as a *process*, involving both the loss and the breaking apart of habitat, habitat fragmentation independent of habitat loss constitutes a difference in *pattern* (or configuration) between landscapes (McGarigal & Cushman 2002). For a given amount of habitat, a more fragmented landscape has more, smaller habitat patches and contains a greater total length of habitat edge (Figure 2). For the remainder of this paper, when I use the term 'habitat fragmentation,' I specifically mean habitat fragmentation independent of habitat loss (i.e. habitat fragmentation *per se*).

Importantly, studies of patch size effects and patch isolation effects do not provide evidence for effects of habitat fragmentation, because both patch size effects and patch isolation effects are inherently confounded with effects of habitat amount (Fahrig 2003; 2013). Smaller patches have less habitat than larger patches, and more isolated patches are more isolated precisely because there is less habitat surrounding them than surrounding less isolated patches (Moilanen & Nieminen 2002; Bender et al. 2003; Tischendorf et al. 2003; Prugh, 2009; Ranius et al. 2010; Thornton et al. 2011; Martin and Fahrig 2012; Figure 3). Therefore, effects of habitat fragmentation must be measured at the landscape scale (McGarigal & Cushman 2002; Figure 4), that is, by comparing effects of habitat pattern across multiple landscapes, while controlling for the total amount of habitat in them.

Researchers have used three different ways to estimate ecological responses to habitat fragmentation. The first is to conduct a true landscape-scale experiment in which one constructs multiple landscapes either across independent gradients of habitat amount and fragmentation (e.g. With et al. 2002; Figure 5) or across a gradient of fragmentation where all constructed landscapes have the same amount of habitat (e.g. Wolff et al. 1997; Goodwin & Fahrig 2002). The second approach for estimating effects of habitat fragmentation is to select a set of existing landscapes across gradients in habitat loss and fragmentation. The landscapes may be selected either randomly (Smith et al. 2011) or in an attempt to minimize any correlation between habitat amount and fragmentation (e.g. Hovel & Lipcius 2001; Cushman & McGarigal 2003; Robertson et al. 2013; Plecaš et al. 2014). A variety of statistical methods are then used to estimate the effects of habitat fragmentation while statistically controlling for the effects of habitat amount. In the third approach, SLOSS ('single large or several small') studies compare species richness across virtual landscapes - sets of patches that are subsets of a larger set, each containing the same total amount of habitat but varying in fragmentation (Figures 6, 7).

3. REVIEW OBJECTIVES AND CRITERIA

In an earlier review (Fahrig 2003) I compared the effects of habitat fragmentation to the effects of habitat loss on a wide range of ecological responses and found that in general the effects of fragmentation are much weaker than the effects of habitat loss; usually no significant response to fragmentation is detected. In that review I also noted that, of the 31 cases in which studies did detect a significant response to habitat fragmentation, only 10 of these responses were negative while the remaining 21 were positive. Given the small sample size at that time (31 significant responses), it would have been premature to conclude that significant responses to habitat fragmentation are predominantly positive.

The current review addresses the questions: (1) are most significant ecological responses to habitat fragmentation negative or positive?, and (2) do particular attributes of species or landscapes lead to a predominance of negative or positive significant responses to habitat fragmentation? I attempted a complete search for studies documenting significant responses to habitat fragmentation, whether positive or negative. Included studies used one of the three landscape-scale study designs described above to control for effects of habitat amount while estimating effects of habitat fragmentation: (i) multi-landscape fragmentation experiment, (ii) observational study of multiple landscapes with statistical control of habitat amount effects, and (iii) SLOSS-type analysis.

I used the following criteria to screen papers for inclusion in this review.

- 1. The significant response to habitat fragmentation was based directly on an empirical test.
- 2. The ecological response variable was directly measured, not inferred e.g. from a model.
- 3. The study included multiple landscapes, varying in degree of fragmentation.
- 4. The study evaluated responses to fragmentation of habitat or natural cover types such as forest, wetland, grassland, coral reef, etc. The response variable (species abundance, richness, etc.) was measured within the habitat or natural cover.
- 5. The study used landscape-scale (not patch-scale) measure(s) of habitat fragmentation; these could be either 'class-level' or 'landscape-level' measures (*sensu* McGarigal 2015).
- 6. The effect of habitat fragmentation was estimated independently of the effect of habitat amount, using one of the three study designs described above.
- 7. The response variable was one for which a direction of fragmentation effect (negative or positive) could be assigned, e.g., abundance, species richness, movement success, etc. This excluded, for example, effects on community composition (e.g., Cisneros et al. 2015) or effects on community mean trait values (e.g., Perović et al. 2015).
- 8. The effect of fragmentation was statistically significant (negative or positive; α=0.05), or fragmentation was in the top model set (in multi-model inference), or fragmentation had a high probability (≥ 0.7) of a non-zero coefficient; for simplicity, I refer to all of these as 'significant effects of fragmentation' below. For SLOSS-type studies I defined a significant effect as one where the largest-to-smallest and smallest-to-largest curves did not cross (Figure 7).
- 9. Finally, the direction of the significant fragmentation effect had to be non-ambiguous. This resulted in exclusion of a few cases in which the effect of fragmentation was significant, but it was non-monotonic or the direction varied over time and/or over spatial scales (e.g. Alofs et al. 2014; With 2015).

To find potential papers I searched for all papers published up to the end of 2015, using the following search term sequence in Web of Science: "fragmentation per se" or SLOSS or (("edge density" or "edge length" or "number of patches" or "mean patch size" or "boundary length" or "patch density" or "median patch size" or "clumping index" or "splitting index" or "aggregation index" or "like adjacencies" or "fractal dimension" or "IJI" or "mean circumscribing circle" or "largest patch index" or "shape index" or "mean core area" or "proportion core area" or "mean nearest-neighbour" or "mean perimeter to area" or "mean edge to area") and (habitat or forest or grassland or wetland or coral or landscapes or watersheds or catchments)). This procedure resulted in over 5,000 'hits'. Most of these were eliminated based on the titles, commonly because the study did not include an ecological response. Based on the abstracts, most of the remainder were eliminated for failure to meet one or more of the criteria above. In many cases suitability could not be determined based on the abstract, so some additional papers were

eliminated after I read the methods and results sections. I also discovered additional papers in the literature cited sections of a few of the included papers.

I took the significant effects of fragmentation from the tables, figures and appendices of the papers. I did not take significant effects from the text portions of the papers, in an effort to reduce bias caused by the tendency of authors to highlight results that support their preconceived notions (Fahrig *in press*). This is a well-known phenomenon, termed 'theory tenacity' or 'confirmation/expectation bias' (Loehle 1987; Nickerson 1998; Jeng 2006). I did not calculate any effects, and I recorded only the sign - positive or negative - of only the significant effects. I did not attempt to estimate the overall effect size of habitat fragmentation because that would have required also extracting many hundreds of non-significant effects of fragmentation; see Section 6, 'Limitations and caveats' below. In addition, by limiting the review to the significant effects, I included the studies with sufficient statistical power to detect a significant effect.

4. MOST SIGNIFICANT RESPONSES TO HABITAT FRAGMENTATION ARE POSITIVE

I found 381 significant responses to habitat fragmentation independent of habitat amount effects, drawn from 118 studies. 290 (76%) of the significant fragmentation effects were positive (Figure 8).

Most significant responses to fragmentation were positive, irrespective of the method used to control for habitat amount effects - experimental, statistical, or SLOSS-type control (Figure 9a). All 53 significant SLOSS-type responses indicated higher species richness in several small patches than in a single large patch, i.e., significant positive responses to fragmentation. This same result was found in early reviews of empirical SLOSS-type studies by Simberloff & Abele (1982) and by Quinn & Harrison (1988). Despite this, authors of SLOSS-type studies published after Quinn and Harrison (1988) are still surprised when they find positive effects of fragmentation. For example, Tsharntke et al. (2002) refer to their "surprising observation that patchy habitat supported more butterfly species than did the same area composed of only one or two fragments, even when only endangered species were considered." The fact that authors continue to be surprised by significant positive effects of fragmentation indicates the level of entrenchment of the current paradigm that assumes negative effects of habitat fragmentation.

Most significant responses to fragmentation were positive, irrespective of the taxonomic group studied (Figure 9b). Birds showed a slightly higher proportion of significant negative responses (36%) than other taxa. I speculate that this is at least partly due to a bias in bird studies, as some authors specifically selected study species that were thought *a priori* to be sensitive to fragmentation (e.g., Smith et al. 2011).

Most significant responses to fragmentation were also positive irrespective of whether fragmentation was measured as number of patches, edge density, mean patch size, or as a whole-landscape ('landscape-level' *sensu* McGarigal 2015) metric (Figure 9c). In contrast, for studies using a 'clumping' metric of fragmentation, 23 of the 41 (56%) significant responses to habitat fragmentation were negative (i.e. positive effects of clumping). However, I note that 11 of these 23 significant negative effects were taken from a single study (Radford & Bennett 2007), in which some individual species showed both significant negative responses to fragmentation measured as clumping and significant positive responses to fragmentation using other metrics of fragmentation. On closer inspection of that study it appears that the application of the clumping metric in the data analysis introduced an unintended positive association between clumping and habitat amount (Figure 10). Therefore, in that study it is possible that the significant positive

effects of clumping (negative effects of fragmentation) may be due to a positive effect of habitat amount.

Most significant responses to habitat fragmentation were also positive regardless of whether the ecological response was a single-species or a multi-species response (generally species richness) (Figure 9d). Of the 231 single-species significant effects, 13 were on movement success and 9 were on demographic responses such as survival rate or population growth rate. Most of these significant effects of habitat fragmentation were also positive (Figure 9d).

4.1. Responses of Habitat Specialists and Threatened or Declining Species

Within the significant species richness responses, several authors evaluated effects of fragmentation on species richness of habitat specialists and/or rare and/or threatened species (30 significant responses). Often this was done in response to the author's surprise at finding significant positive effects of habitat fragmentation on total species richness. Authors reasoned that perhaps in fragmented landscapes, habitat patches contain more generalist species or even matrix-preferring species. The higher habitat edge length in more fragmented landscapes would then result in more 'spillover' of such species into habitat patches in more fragmented landscapes. This suggestion was not supported by the data. Twenty-nine (97%) of the 30 significant responses of specialist/rare/threatened species richness to fragmentation were positive (Figure 9d).

Similarly, for single-species responses, most significant effects of fragmentation were positive, irrespective of the IUCN status (IUCN 2015) of the species (Figure 9e). The majority of significant single-species responses were positive, irrespective of whether the species is listed as increasing, stable, or declining in abundance. Two of the species showing significant responses to habitat fragmentation are listed as endangered; one showed a negative response and the other a positive response to fragmentation. An additional five species are listed as near-threatened; all five showed significant positive responses to habitat fragmentation.

4.2 Tests of Hypotheses Predicting Negative Fragmentation Effects

In the current paradigm, which assumes that effects of fragmentation are generally negative, several hypotheses have been suggested to predict situations in which negative effects of fragmentation should be particularly likely. First, some authors have suggested that negative effects of forest fragmentation are more likely in tropical than in temperate regions, because forest animals in the tropics may be less likely to cross open areas than forest animals in temperate regions (Báldi 1996; Lindell et al. 2007; Cerezo et al. 2010). I found no support for this hypothesis. The majority of significant responses to fragmentation were positive, irrespective of biome (Figure 11a).

Second, it has been suggested that species that interact with the landscape over larger areas are more likely to show negative effects of habitat fragmentation, as they are more likely to encounter non-habitat ('matrix') areas during their movements. I could not directly evaluate this idea because reliable movement range data are not available for the wide range of species and species groups covered in this review. However, I reasoned that the sizes of the replicate landscapes chosen for a study might be a rough indicator of the movement range of the species (Jackson & Fahrig 2012) and species groups in it, as authors should select the sizes of replicate landscapes to reflect movement range. Sizes of replicate landscapes varied across studies by four

orders of magnitude. However, the majority of significant fragmentation effects were positive, irrespective of the sizes of replicate landscapes (Figure 11b). Therefore, the review did not support the movement range hypothesis. It seems likely that the expectation of negative effects of fragmentation on species with larger movement ranges is related more to effects of habitat loss than to effects of habitat fragmentation. Species with large movement ranges may have high habitat requirements, but that does not mean their habitat must be contiguous.

Finally, several theoretical studies have suggested that negative responses to habitat fragmentation should be more likely when habitat amount is low, particularly when habitat covers less than about 20-30% of the landscape, the so-called 'fragmentation threshold' (Bascompte & Solé 1996; Boswell et al. 1998; Fahrig 1998; Flather & Bevers 2002; Hill & Caswell 1999; Lande 1987; With & King 1999). However, I found no evidence for the fragmentation threshold hypothesis in this review; the majority of significant responses to habitat fragmentation were positive, irrespective of the median level of habitat amount in the study (Figure 11c). Interestingly, With (2015) did find that fragmentation effects are more likely at low habitat amount (10-20%) than high habitat amount (60-80%); however, the response to fragmentation at low habitat amount was positive.

5. AUTHORS' EXPLANATIONS FOR SIGNIFICANT POSITIVE RESPONSES TO HABITAT FRAGMENTATION

According to the Introduction sections of papers, most authors had *a priori* expected negative responses to habitat fragmentation, based on one (or more) of three assumptions: (i) negative edge effects, especially due to higher assumed predation or nest parasitism at habitat edges, (ii) lower connectivity in landscapes with many small patches than in landscapes with a few large patches, and (iii) minimum patch size effects. In the Discussion sections of papers in which authors report significant negative effects of habitat fragmentation, they generally refer to these effects as "expected." Thus, the reigning paradigm assumes negative effects of habitat fragmentation.

The results of this review run counter to this paradigm, with 76% of significant responses to habitat fragmentation positive. When authors obtained a significant positive response to fragmentation, they often expressed surprise in the Discussion section, e.g. "we obtained puzzling results ... fragmented populations had larger abundances than nonfragmented populations" (Bancroft & Turchin 2003); "our results do not support the underlying theoretical supposition that regional extinction should increase with increasing degrees of habitat subdivision" (Robinson & Quinn 1988).

Given the unexpected preponderance of positive significant responses to fragmentation, I scoured the Discussion sections of the studies containing these positive responses to extract the authors' explanations for them (Figure 12). Surprisingly, authors offered no explanations for 119 (41%) of the significant positive responses. While I have no way of knowing the reason for this omission I speculate that many authors are uncomfortable accepting the notion that their data indicate positive responses to fragmentation. In addition to the 41% of cases in which no explanation was provided, in 10 cases the authors offered an explanation, but I was unable to understand it as an explanation for a positive response to fragmentation. The following review is therefore based on authors' explanations for the remaining 161 significant positive fragmentation effects.

5.1. Fragmentation Increases Functional Connectivity

The most common explanation offered by authors for significant positive responses to habitat fragmentation was that it increases functional connectivity of the landscape. That fragmentation can increase landscape functional connectivity is supported by the fact that 9 of the 13 significant effects of fragmentation on movement success were positive (Figure 9d).

Authors suggested two main mechanisms by which fragmentation can increase functional connectivity. First, higher fragmentation implies a larger number of smaller patches, with smaller distances between them (Figure 2). This topology can increase patch encounter rate, thus increasing patch immigration and reducing emigration from the system of patches. For example, "[h]igh faunal densities in patchy seagrass has (sic) been linked to high perimeter-to-area ratios of small patches, which may increase the probability that dispersing organisms will encounter the patch" (Healey & Hovel 2004). This mechanism for a positive influence of habitat patchiness on movement success has also been shown in theoretical studies (Tischendorf & Fahrig 2000; Bowman et al. 2002; Saura et al. 2014).

The second mechanism suggested by authors for higher functional connectivity with higher fragmentation was that increasing edge density facilitates movement for species that preferentially move along edges. For example, "the positive effect of edge density … suggests that field edges provided connectivity and facilitated wasp movements between trap nests and source habitats where dispersal started" (Holzschuh et al. 2010) and, "it may be that functional connectivity is higher in landscapes with a linear network of riparian and roadside vegetation" (Radford et al. 2005).

5.2. More Fragmented Landscapes Contain a Higher Diversity of Habitats

Among papers measuring effects of habitat fragmentation on species richness, e.g., SLOSS- type analyses, habitat diversity was the most common explanation offered by authors for significant positive responses to fragmentation. For example, "a series of small- or medium-sized reserves capture a much greater habitat and habitat—environment heterogeneity than one large fragment" (Tscharntke et al. 2002) or, "a large number of patches can provide high habitat diversity for species with different environmental requirements and result in decreased regional extinction risks" (Hu et al. 2012).

5.3 Positive Edge Effects

Authors attributed 32 of the significant positive responses to habitat fragmentation to positive edge effects. They argued that edges are more productive and more structurally diverse than habitat interior, offering higher forage availability and refuge from predation (Walter 2009; Klingbeil & Willig 2009; Henden et al. 2011; Moore et al. 2011). The idea that edges are high-quality habitats for many species goes back to the early 1900s (Leopold 1933).

The suggestion by some authors that habitat fragmentation reduces predation pressure at edges may seem surprising. Some studies have documented higher predation rates at forest edges than in forest interiors, particularly in agricultural regions where important predators are openhabitat generalist species (Robinson et al. 1995), although this is by no means a general pattern (Tewksbury et al. 1998; Carlson & Hartman 2001). Therefore it seems likely that at least some of the 24% significant negative responses to habitat fragmentation are due to negative edge

effects. Negative edge effects are, implicitly, the favoured interpretation for significant negative effects of habitat fragmentation. Many authors made general reference to interior specialist species, edge avoiders, and negative edge effects, particularly in the Introduction sections of papers (e.g. McAlpine et al. 2006; Grossman et al. 2008; Cerezo et al. 2010). Oddly however, I did not find any examples in Discussion sections of authors linking their findings of significant negative effects of fragmentation to evidence of edge avoidance or negative edge effects for *their study species*, whether from their own work or from that of others.

5.4. Increased Persistence of Predator-prey and Host-parasitoid Systems

Authors attributed 23 of the significant positive effects of habitat fragmentation to the idea that habitat fragmentation stabilizes or increases persistence of predator-prey and host-parasitoid systems. As mentioned above in Section 1. 'Habitat patchiness', this was the earliest explanation for positive effects of habitat patchiness. Fragmentation is thought to increase persistence of predator-prey and host-parasitoid systems either by offering refuges for the prey or host or because it reduces the dispersal efficiency of the predator or parasitoid, allowing the prey to stay a step ahead of the predator in space and time; e.g. "fragmented boreal forest may limit dispersal of parasitoids" (Roland 1993).

5.5. Habitat Fragmentation Reduces Intra-specific and Inter-specific Competition

Authors attributed 19 significant positive effects of fragmentation to negative effects of fragmentation on intra-specific or inter-specific competition. For example, authors suggest that fragmentation allows more small mammal territories (Wolff et al. 1997; Collins & Barrett 1997) and more crab territories (Caley et al. 2001) to be defended, because the edges of individual patches are used as territory boundaries. In addition, theory suggests that fragmentation can increase coexistence of species if the stronger competitors are the weaker dispersers (Levins & Culver 1971), and some authors invoked this mechanism. For example, Dufour et al. (2006) argued that "spatial structure may indefinitely delay competitive exclusion" by allowing "repeated immigration" of poor competitors, and Hanski (1987) described decreasing covariance of competing species with increasing habitat fragmentation.

5.6. Habitat Fragmentation Spreads the Risk of Extinction

Seven significant positive responses to habitat fragmentation were attributed by authors to the idea that habitat fragmentation spreads the risk of extinction over a larger number of sites, thus reducing the risk of simultaneous extinction of all local populations (den Boer 1968). For example, Martínez-Sanz et al. (2012) argued that, "the presence of a number of ponds relatively close to each other and interconnected by dispersal can be particularly important for biodiversity." Note that the spreading of risk explanation is a general form of the previous two explanations, as "risk" can be caused not only by disturbances but also by predation, parasitism, or competition.

5.7. Habitat Fragmentation Increases Landscape Complementation

Finally, a few authors argued that, for species that require two or more different kinds of habitat for persistence, habitat fragmentation increases accessibility of these different habitat types, i.e., it increases 'landscape complementation' *sensu* Dunning et al. (1992). When a habitat type is more fragmented, it is more interspersed with other required habitat type(s), increasing access to them (Figure 13). As argued by Slancarova et al. (2014), "A landscape with heterogeneous configuration contains multiple edges and notches, which support species requiring diverse resources located in close proximity."

5.8. Which Explanation is Correct?

At this point it is not possible to draw conclusions as to which of the mechanism(s) causing significant positive fragmentation effects are most important. In most cases the explanation provided by the authors was *post hoc* speculation rather than an *a priori* prediction because authors often had assumed, before conducting the study, that any detected responses to habitat fragmentation would be negative. In addition, a given author was likely to invoke the same explanation for most or all significant positive fragmentation effects found in his/her study, which likely over-weights some explanations. Some authors suggested the combined actions of more than one mechanism. Overall, it seems likely that all of the mechanisms above, and possibly additional ones, operate in at least some situations.

5.9. Matrix Quality and Landscape Heterogeneity

An additional possible explanation for significant positive effects of habitat fragmentation is that the matrix or non-habitat portion of the landscape may often be more wildlife-friendly in landscapes containing many small habitat patches than in landscapes containing a few large habitat patches. This seems particularly likely for agricultural landscapes where crop fields form the matrix. In fact, several studies have found positive relationships between landscape configurational heterogeneity (*sensu* Fahrig & Nuttle 2005; Fahrig et al. 2011) and biodiversity in agricultural landscapes (Concepción et al. 2012; Flick et al. 2012; Lindsay et al. 2013; Fahrig et al. 2015).

Of course the opposite could be true in some cases, and this may explain some of the significant negative responses to habitat fragmentation. For example, in a particular region, if landscapes with many small patches contain more roads than landscapes with a smaller number of large patches, the matrix quality of the former will be lower than the matrix quality of the latter, which could lead to higher road mortality in the former and thus an apparent negative response to habitat fragmentation. Elucidation of the possible role of matrix quality in creating positive or negative significant responses to habitat fragmentation will require studies in which variables such as road density, cropping system, etc. are included along with the habitat amount and fragmentation variables.

The potential role of matrix quality in mediating significant positive or negative responses to habitat fragmentation suggests a more holistic view of the landscape may be needed. The idea that patchiness or spatial heterogeneity of the whole landscape can benefit biodiversity was originally proposed by Duelli (1997) as the "mosaic concept." Duelli argued that important positive correlates of biodiversity are the amount of natural and semi-natural area,

the number of patches, and the length of ecotones between patch types. Duelli explicitly recognized that significant positive effects of landscape spatial heterogeneity run counter to habitat patch size and isolation effects as inferred from the theory of island biogeography (see Figure 6 in Duelli 1997). While Duelli discussed his mosaic concept only in the context of agricultural landscapes, the results of this review suggest that positive effects of landscape spatial heterogeneity may be more general. For example, Sattler et al. (2010) found increasing arthropod diversity with increasing landscape spatial heterogeneity within cities in Switzerland. Hovick et al. (2015) created replicate grassland landscapes on a gradient of increasing landscape spatial heterogeneity and found positive relationships between landscape heterogeneity and bird diversity and community stability.

6. LIMITATIONS AND CAVEATS

The overall conclusion of this review, that most significant responses to habitat fragmentation are positive, appears to be robust to a wide variety of conditions. I attempted to identify sets of conditions that differentiate between mostly positive and mostly negative significant fragmentation effects, but I did not find any situations in which most effects were negative. However, I note that this failure may be an artefact of the structure of the data. The conditions represented by the 381 significant responses - taxa, study types, response and predictor variables, spatial scales, etc. - are not mutually independent. For example, SLOSS-type studies all used the same response (species richness), and studies of birds used almost exclusively single-species responses. Thus, the fact that I did not find any conditions in which most significant responses were negative does not mean that such situations do not exist. I reiterate, though, that this review shows that across a very wide range of conditions the majority of significant responses to habitat fragmentation are positive.

It is also important to bear in mind that the first aim of this review was to answer the question, "Are most significant responses to habitat fragmentation negative or positive?" Therefore, I included only the *significant* effects of habitat fragmentation. This review does not answer the question, "What is the overall effect of habitat fragmentation?" As I discuss elsewhere (Fahrig 2003; 2013), the overall effect of habitat fragmentation appears to be very weak. As this review does not include papers finding no significant effects of habitat fragmentation, I do not know the number of such non-significant effects. However, I attempted a rough estimate using the mean proportion of effects of fragmentation that are significant, across the studies in my review containing at least 10 responses, typically responses of different species. For this estimate I excluded studies that tested effects of multiple fragmentation metrics on each response, as this would inflate Type I error. Fourteen of the studies in my review met these criteria. Across these 14 studies the mean proportion of responses to habitat fragmentation that were significant was 0.29 (s.d. 0.22). Thus, a rough estimate of the number of non-significant effects of fragmentation is 381/0.29 or about 1300 non-significant effects. The true number is likely higher, as the 14 studies used to make this estimate are biased, in that each of them had at least one significant effect of habitat fragmentation, a criterion for inclusion in my review. Thus, with a predominance of non-significant effects, and the fact that significant effects are both negative and positive, there is likely very little overall effect of habitat fragmentation on ecological responses (see also Fahrig 2003, 2013).

It is also important to keep in mind that, in this review, when I refer to a significant "positive" relationship, I am referring only to the direction of the relationship. No value

judgment is intended. In some cases a positive relationship may be viewed as a negative effect, e.g., in the case of invasive species, pest species, or over-abundant species. Such value judgments can differ depending on the geographic location. Similarly, in some situations species diversity may be less desirable than a system state with lower species diversity. However, the majority of significant responses to habitat fragmentation were positive even for threatened/specialist species, and irrespective of IUCN status. Therefore, applying value judgments to the definition of "positive" would not change the overall conclusion of this review.

7. FINAL CONSIDERATIONS

7.1 The Fragmentation Zombie

Zombie ideas are ideas that should be dead but are not (Fox 2011). The results of this review suggest that the idea that habitat fragmentation, independent of habitat loss, has widespread negative effects on ecological responses qualifies as a zombie idea. It arose from (i) confounding habitat patchiness with habitat loss (see Section 1) and (ii) inappropriate extrapolation of patch-scale patterns to landscape-scale inferences. The fact that this zombie has persisted for over 45 years is a testament to its intuitive appeal (Fahrig *in press*).

This review also suggests that the fragmentation zombie actually encompasses a whole family of zombie ideas. For example, the general expectation that a small number of large patches should contain more species than a large number of small patches persists, despite consistent reviews showing the opposite (Simberloff & Abele 1982; Quinn & Harrison 1988; this paper). Other members of the fragmentation zombie family include the ideas that: edge effects are generally negative; fragmentation reduces connectivity; habitat specialists show particularly negative responses to fragmentation; and negative fragmentation effects are particularly strong at low levels of habitat amount and in the tropics. The results of this review challenge all of these long-standing, persistent ideas.

7.2 Reframing the fragmentation message

It is important to be very clear about the fragmentation message implied by the results of this review. If, as shown here, significant effects of habitat fragmentation are more often positive than negative, this implies the need for a *more* cautious approach to habitat conservation. Significant positive responses to fragmentation mean there is no justification for assigning lower conservation value to a small patch than to an equivalent area within a large patch, just the opposite. The results also suggest that, generally speaking, land-sharing policies will provide higher ecological value than land-sparing policies (Figure 2).

It is also important to keep in mind that about 24% of significant responses to habitat fragmentation are negative. Therefore, there are likely some situations in which minimizing habitat fragmentation is a reasonable conservation goal. This might occur when conservation action is aimed at a particular species that is known to show a negative response to habitat fragmentation. However, I emphasize that the results of this review do not suggest more significant negative effects of fragmentation for more threatened species; if anything, they suggest the opposite (Figure 9d, e). Thus, while fragmentation has significant negative effects in some particular situations, these cannot be generalized.

The results of this review also indicate that the language associated with much of the habitat loss and fragmentation literature is leading to erroneous conclusions. Prior to about 2005, it was common to see statements such as, "habitat fragmentation is a major cause of biodiversity erosion" (Tabarelli et al. 1999) or, "habitat fragmentation is a leading cause of extinction" (Bruna & Oli 2005). These statements clearly confounded habitat fragmentation with habitat loss. They summarized the patch-scale findings on patch size and isolation effects which, as discussed above (Section 2. "How to measure effects of habitat fragmentation"), are strongly confounded with effects of habitat loss and do not provide information on independent effects of habitat fragmentation.

Since about 2005, many authors acknowledge that habitat loss and fragmentation are separate, the former being a process and the latter being a pattern (Figure 2). However, most authors still assume that the effects of habitat fragmentation independent of the effects of habitat loss are generally negative, as evidenced in the following statements, "habitat loss and fragmentation are major threats to terrestrial biodiversity" (Prugh et al. 2008), "habitat loss and fragmentation are the principal causes of the loss of biological diversity" (Mbora & McPeek 2009), "habitat loss and fragmentation cause significant loss of species richness" (Barth et al 2015), and "habitat loss and fragmentation inevitably cause biodiversity decline" (Barelli et al. 2015).

The results of this review indicate that such statements are in fact false. While habitat loss is, without doubt, one of the most significant causes of biodiversity decline, the significant responses to habitat fragmentation independent of habitat amount are rare, and mostly positive.

ACKNOWLEDGMENTS

I am grateful for helpful comments on previous versions of this paper and related presentations from: Igor Coelho, Sara Collins, Susie Crowe, Michelle Fairbrother, Sandra Martins de Freitas, Jessica Gurevitch, Jochen Jaeger, Jean-Louis Martin, Andrew Moraga, Anne Munier, Genevieve Perkins, Margaret Sawatsky, Chris Souliere, Lutz Tischendorf, and members of the Geomatics and Landscape Ecology Friday Discussion Group. This work was supported by the Natural Sciences and Engineering Research Council of Canada (NSERC).

LITERATURE CITED

- Abele LG, Patton WK. 1976. The size of coral heads and the community biology of associated decapod crustaceans. *Journal of Biogeography* 3:35-47
- Aben J, Adriaensen F, Thijs KW, Pellikka P, Siljander M, Lens L, Matthysen E. 2012. Effects of matrix composition and configuration on forest bird movements in a fragmented Afromontane biodiversity hotspot. *Animal Conservation* 15: 658–668
- Alofs KM, Fowler NL. 2010. Habitat fragmentation caused by woody plant encroachment inhibits the spread of an invasive grass. *Journal of Applied Ecology* 47:338-347
- Alofs KM, González AV, Fowler NL. 2014. Local native plant diversity responds to habitat loss and fragmentation over different time spans and spatial scales. *Plant Ecology* 215:1139-1151
- Amiri BJ, Nakane K. 2009. Modeling the linkage between river water quality and landscape metrics in the Chugoku district of Japan. *Water Resources Management* 23:931-956

- Andreassen HP, Ims RA. 2001. Dispersal in patchy vole populations: Role of patch configuration, density dependence, and demography. *Ecology* 82: 2911–2926
- Anzures-Dadda A, Manson RH. 2007. Patch- and landscape-scale effects on howler monkey distribution and abundance in rainforest fragments. *Animal Conservation* 10: 69–76
- Arryo-Rodriguez V, González-Perez IM, Garmendia A, Solà M, Estrada A. 2013. The relative impact of forest patch and landscape attributes on black howler monkey populations in the fragmented Lacandona rainforest, Mexico. *Landscape Ecology* 28:1717-1727
- Atauri JA, de Lucio JV. 2001. The role of landscape structure in species richness distribution of birds, amphibians, reptiles and lepidopterans in Mediterranean landscapes. *Landscape Ecology* 16:147-159
- Báldi A. 1996. Edge effects in tropical versus temperate forest bird communities: Three alternative hypotheses for the explanation of differences. *Acta Zoologica Academiae Scientiarum Hungaricae* 42:163–172
- Báldi A, Kisbenedek T. 2000. Bird species numbers in an archipelago of reeds at Lake Velence, Hungary. *Global Ecology & Biogeography* 9:451-461
- Bancroft JS, Turchin P. 2003. An experimental test of fragmentation and loss of habitat with *Oryzaephilus surinamensis*. *Ecology* 84:1756-1767
- Barelli C, Rovero F, Hodges K, Araldi A, Heistermann M. 2015. Physiological stress levels in the endemic and endangered Udzungwa red colobus vary with elevation. *African Zoology* 50:23-30.
- Barth BJ, FitzGibbon SI, Wilson RS. 2015. New urban developments that retain more remnant trees have greater bird diversity. *Landscape and Urban Planning* 136:122-129
- Bartolommei P, Mortelliti A, Pezzo F, Puglisi L. 2013. Distribution of nocturnal birds (Strigiformes and Caprimulgidae) in relation to land-use types, extent and configuration in agricultural landscapes of Central Italy. *Rendiconti Lincei* 24:13-21
- Bascompte J, Solé RV. 1996. Habitat fragmentation and extinction thresholds in spatially explicit models. *Journal of Animal Ecology* 65:465-473
- Baz A, Garcia-Boyero A. 1996. The SLOSS dilemma: A butterfly case study. *Biodiversity and Conservation* 5:493-502
- Bender DJ, Tischendorf L, Fahrig L. 2003. Using patch isolation metrics to predict animal movement in binary landscapes. *Landscape Ecology* 18:17–39
- Blake JG, Karr JR. 1984. Species composition of bird communities and the conservation benefit of large versus small forests. *Biological Conservation* 30:173-187
- Boscolo D, Metzger JP. 2009. Is bird incidence in Atlantic forest fragments influenced by landscape patterns at multiple scales? *Landscape Ecology* 24:907-918
- Boswell GP, Britton NF, Franks NR. 1998. Habitat fragmentation, percolation theory and the conservation of a keystone species. *Proceedings of the Royal Society of London Series B* 265:1921-1925
- Bowman J, Cappuccino N, Fahrig L. 2002. Patch size and population density: the effect of immigration behavior. *Conservation Ecology*. 6:art.9
- Bruna EM, Oli MK. 2005. Demographic effects of habitat fragmentation on a tropical herb: life-table response experiments. *Ecology* 86:1816–1824
- Caley JM, Buckley KA, Jones GP. 2001. Separating ecological effects of habitat fragmentation, degradation, and loss on coral commensals. *Ecology* 82:3435–3448

- Caravaggi A, Montgomery WI, Reid N. 2015. Range expansion and comparative habitat use of insular, congeneric lagomorphs: invasive European hares *Lepus europaeus* and endemic Irish hares *Lepus timidus hibernicus*. *Biological Invasions* 17:687-698
- Carlson A, Hartman G. 2001. Tropical forest fragmentation and nest predation An experimental study in an Eastern Arc montane forest, Tanzania. *Biodiversity and Conservation* 10:1077-1085
- Carrara E, Arroyo-Rodríguez V, Vega-Rivera JH, Schondube JE, de Freitas SM, Fahrig L. 2015. Impact of landscape composition and configuration on forest specialist and generalist bird species in the fragmented Lacandona rainforest, Mexico. *Biological Conservation* 184:117–126
- Cerezo A, Perelman S, Robbins CS. 2010. Landscape-level impact of tropical forest loss and fragmentation on bird occurrence in eastern Guatemala. *Ecological Modelling* 221:512-526
- Cisneros LM, Fagan ME, Willig MR. 2015. Season-specific and guild-specific effects of anthropogenic landscape modification on metacommunity structure of tropical bats. *Journal of Animal Ecology* 84:373-385
- Clare JDJ, Anderson EM, MacFarland DM. 2015. Predicting bobcat abundance at a landscape scale and evaluating occupancy as a density index in Central Wisconsin. *Journal of Wildlife Management* 79:469-480
- Collinge SK, Forman RTT. 1998. A conceptual model of land conversion processes: predictions and evidence from a microlandscape experiment with grassland insects. *Oikos* 82:66-84
- Collins RJ, Barrett GW. 1997. Effects of habitat fragmentation on meadow vole (*Microtus pennsylvanicus*) population dynamics in experiment landscape patches. *Landscape Ecology* 12:63-76
- Concepción ED, Díaz M, Kleijn D, Báldi A, Batáry P, Clough Y, Gabriel D, Herzog F, Holzschuh A, Knop E, Marshall EJP, Tscharntke T, Verhulst J. 2012. Interactive effects of landscape context constrain the effectiveness of local agri-environmental management. *Journal of Applied Ecology* 49:695-705
- Cooper CB, Walters JR. 2002. Independent effects of woodland loss and fragmentation on Brown Treecreeper distribution. *Biological Conservation* 105:1-10
- Cooper JK, Li J, Montagnes DJS. 2012. Intermediate fragmentation per se provides stable predator-prey metapopulation dynamics. *Ecology Letters* 15:856-863
- Cushman SA, McGarigal K. 2003. Landscape-level patterns of avian diversity in the Oregon Coast Range. *Ecological Monographs* 73:259-281
- Dauber J, Bengtsson J, Lenoir L. 2006. Evaluating effects of habitat loss and land-use continuity on ant species richness in seminatural grassland remnants. *Conservation Biology* 20:1150–1160
- den Boer PJ. 1968. Spreading of risk and stabilization of animal numbers. *Acta Biotheoretica* 18:165-194
- Didham RK, Kapos V, Ewers RM. 2012. Rethinking the conceptual foundations of habitat fragmentation research. *Oikos* 121:161-170
- Dormann CF, Schweiger O, Augenstein I, Bailey D, Billeter R, de Blust G, DeFilippi R, Frenzel M, Hendrickx F, Herzog F, Klotz S, Liira J, Maelfait J-P, Schmidt T, Speelmans M, van Wingerden WKRE, Zobel M. 2007. Effects of landscape structure and land-use intensity on similarity of plant and animal communities. *Global Ecology and Biogeography* 16:774-787

- Duelli P. 1997. Biodiversity evaluation in agricultural landscapes: An approach at two different scales. *Agriculture, Ecosystems and Environment* 62:81-91
- Dufour A, Gadallah F, Wagner HH, Guisan A, Buttler A. 2006. Plant species richness and environmental heterogeneity in a mountain landscape: Effects of variability and spatial configuration. *Ecography* 29:573-584
- Dunning JB, Danielson BJ, Pulliam HR. 1992. Ecological processes that affect populations in complex landscapes. *Oikos* 10:169-195
- Dzwonko Z, Loster S. 1989. Distribution of vascular plant species in small woodlands on the Western Carpathian foothills. *Oikos* 56:77-86
- Ethier K, Fahrig L. 2011. Positive effects of forest fragmentation, independent of forest amount, on bat abundance in eastern Ontario, Canada. *Landscape Ecology* 26:865-876
- Fahrig L. 1998. When does fragmentation of breeding habitat affect population survival? *Ecological Modelling* 105:273-292
- Fahrig L. 2003. Effects of habitat fragmentation on biodiversity. *Annual Reviews of Ecology, Evolution and Systematics* 34:487-515
- Fahrig L. 2013. Rethinking patch size and isolation effects: the habitat amount hypothesis. *Journal of Biogeography* 40:1649-1663
- Fahrig L. *in press*. Forty years of bias in habitat fragmentation research. In: Kareiva P, Silliman B, Marvier M. *Uncomfortable questions and confirmation bias in conservation*. Oxford University Press, Oxford, UK.
- Fahrig L, Baudry J, Brotons L, Burel FG, Crist TO, Fuller RJ, Sirami C, Siriwardena GM, Martin J-L. 2011. Functional heterogeneity and biodiversity in agricultural landscapes. *Ecology Letters* 14:101-112.
- Fahrig L, Girard J, Duro D, Pasher J, Smith A, King D, Lindsay K, Mitchell S, Tischendorf L. 2015. Farmlands with smaller crop fields have higher within-field biodiversity. *Agriculture, Ecosystems and Environment* 200:219-234.
- Fahrig L, Nuttle WK. 2005. Population ecology in spatially heterogeneous environments. Pp. 95-118 in: Lovett GM, Jones CG, Turner MG, Weathers KC (eds.). *Ecosystem function in heterogeneous landscapes*. Springer-Verlag, New York.
- Fattorini S. 2010. The use of cumulative area curves in biological conservation: A cautionary note. *Acta Oecologica* 36:255-258
- Flather CH, Bevers M. 2002. Patchy reaction-diffusion and population abundance: the relative importance of habitat amount and arrangement. *American Naturalist* 159:40-56
- Flick T, Feagan S, Fahrig L. 2012. Effects of landscape structure on butterfly species richness and abundance in agricultural landscapes in eastern Ontario, Canada. *Agriculture*, *Ecosystems and Environment* 156:123-133
- Fox J. 2011. Zombie ideas in ecology. *Oikos Blog* https://oikosjournal.wordpress.com/2011/06/17/zombie-ideas-in-ecology/
- Game M, Peterken GF. 1984. Nature reserve selection strategies in the woodlands of Central Lincolnshire, England. *Biological Conservation* 29:157-181
- Gause GF. 1934. *The struggle for existence*. The Williams & Wilkins Company, Baltimore, MD, USA
- Gavish Y, Ziv Y, Rosenzweig ML. 2012. Decoupling fragmentation from habitat loss for spiders in patchy agricultural landscapes. *Conservation Biology* 26:150–159

- Gémesi Z, Downing JA, Cruse RM, Anderson PF. 2011. Effects of watershed configuration and composition on downstream lake water quality. *Journal of Environmental Quality* 40:517–527
- Gilpin ME, Diamond JM. 1980. Subdivision of nature reserves and the maintenance of species diversity. *Nature* 285:567-568
- Gonzalez-Moreno P, Pino J, Carreras D, Basnou C, Fernández-Rebollar I, Vilà M. 2013. Quantifying the landscape influence on plant invasions in Mediterranean coastal habitats. *Landscape Ecology* 28:891-903
- Goodwin BJ, Fahrig L. 2002. How does landscape structure influence landscape connectivity? *Oikos* 99:552-570
- Green RE, Cornell SJ, Scharlemann JPW, Balmford A. 2005. Farming and the fate of wild nature. *Science* 307:550-555
- Grez AA, Zaviezo T, Ríos M. 2005. Ladybird (Coleoptera: Coccinellidae) dispersal in experimental fragmented alfalfa landscapes. European Journal of Entomology 102:209-216
- Grez A, Zaviezo T, Tischendorf L, Fahrig L. 2004. A transient, positive effect of habitat fragmentation on insect population densities. *Oecologia* 141:444-451
- Grossman SR, Hannon SJ, Sánchez-Azofeifa A. 2008. Responses of Great Horned Owls (*Bubo virginianus*), Barred Owls (*Strix varia*), and Northern Saw-whet Owls (*Aegolius acadicus*) to forest cover and configuration in an agricultural landscape in Alberta, Canada. *Canadian Journal of Zoology* 86:1165-1172
- Hagan J M, McKinley PS, Meehan AL, Grove SL. 1997. Diversity and abundance of landbirds in a northeastern industrial forest. *Journal of Wildlife Management* 61:718–735
- Haila Y, Hanski IK. 1984. Methodology for studying the effect of habitat fragmentation on land birds. *Annales Zoologici Fennici* 21: 393-397
- Hanski I. 1987. Carrion fly community dynamics: patchiness, seasonality and coexistence. *Ecological Entomology* 12:257-266

Conservation 18:307-315

- Hanski I. 2015. Habitat fragmentation and species richness. *Journal of Biogeography* 42:989-994 Harms TM, Kinkead KE, Dinsmore SJ. 2014. Evaluating the effects of landscape configuration on site occupancy and movement dynamics of odonates in Iowa. *Journal of Insect*
- Hattori A, Shibuno T. 2010. The effect of patch reef size on fish species richness in a shallow coral reef shore zone where territorial herbivores are abundant. *Ecological Research* 25:457-468
- Healey D, Hovel KA. 2004. Seagrass bed patchiness: effects on epifaunal communities in San Diego Bay, USA. *Journal of Experimental Marine Biology and Ecology* 313:155-174
- Henden J-A, Ims RA, Yoccoz NG, Sørensen R, Killengreen ST. 2011. Population dynamics of tundra voles in relation to configuration of willow thickets in southern arctic tundra. *Polar Biology* 34:533-540
- Henden J-A, Yoccoz NG, Ims RA, Langeland K. 2013. How spatial variation in areal extent and configuration of labile vegetation states affect the riparian bird community in Arctic tundra. *PLOS ONE* 8:e63312
- Hicks JP, Hails RS, Sait SM. 2015. Scale-dependent, contrasting effects of habitat fragmentation on host-natural enemy trophic interactions. *Landscape Ecology* 30: 1371-1385.
- Higgs AJ, Usher MB. 1980. Should nature reserves be large or small? *Nature* 285:568-569
- Hill MF, Caswell H. 1999. Habitat fragmentation and extinction thresholds on fractal landscapes. *Ecology Letters* 2:121-127

- Hiron M, Berg Å, Eggers S, Berggren Å, Josefsson J, Pärt T. 2015. The relationship of bird diversity to crop and non-crop heterogeneity in agricultural landscapes. *Landscape Ecology* 30:2001-2013
- Hokkanen PJ, Kouki J, Komonen A. 2009. Nestedness, SLOSS and conservation networks of boreal herb-rich forests. *Applied Vegetation Science* 12: 295–303
- Holyoak M, Lawlor SP. 1996. The role of dispersal in predator-prey metapopulation dynamics. *Journal of Animal Ecology* 65:640-652
- Holzkamper A, Lausch A, Seppelt R. 2006. Optimizing landscape configuration to enhance habitat suitability for species with contrasting habitat requirements. *Ecological Modelling* 198:277-292
- Holzschuh A, Steffan-Dewenter I, Tscharntke T. 2010. How do landscape composition and configuration, organic farming and fallow strips affect the diversity of bees, wasps and their parasitoids? *Journal of Animal Ecology* 79:491-500
- Hoss SK, Guyer C, Smith LL, Schuett GW. 2010. Multiscale influences of landscape composition and configuration on the spatial ecology of eastern diamond-backed rattlesnakes (*Crotalus adamanteus*). *Journal of Herpetology* 44:110-123
- Hovel KA. 2003. Habitat fragmentation in marine landscapes: relative effects of habitat cover and configuration on juvenile crab survival in California and North Carolina seagrass beds. *Biological Conservation* 110:401-412
- Hovel KA, Lipcius RN. 2001 Habitat fragmentation in a seagrass landscape: Patch size and complexity control blue crap survival. *Ecology* 82:1814-1829
- Hovick TJ, Elmore RD, Fuhlendorf SD, Engle DM, Hamilton RG. 2015. Spatial heterogeneity increases diversity and stability in grassland bird communities. *Ecological Applications* 25:662-672
- Hu G, Wu J, Feeley KJ, Xu G, Yu M. 2012. The effects of landscape variables on the species-area relationship during late-stage habitat fragmentation. *PLOS ONE* 7: e43894
- Huffaker CB. 1958. Experimental studies on predation: Dispersion factors and predator-prey oscillations. *Hilgardia* 27:795-835
- IUCN. 2015. The IUCN red list of threatened species. http://www.iucnredlist.org/
- Jackson, HB, Fahrig L. 2012. What size is a biologically relevant landscape? *Landscape Ecology* 27: 929-941.
- Järvinen O. 1982. Conservation of endangered plant populations: Single large or several small reserves? *Oikos* 38:301-307
- Jeng, M. 2006. A selected history of expectation bias in physics. *American Journal of Physics* 74:578-583
- Johnson LB, Richards C, Host GE, Arthur JW. 1997. Landscape influences on water chemistry in Midwestern stream ecosystems. *Freshwater Biology* 37:193-208
- Kie JG, Bowyer RT, Nicholson MC, Boroski BB, Loft ER. 2002. Landscape heterogeneity at differing scales: Effects on spatial distribution of mule deer. *Ecology* 83:530-544
- King D I, Griffin CR, DeGraaf RM. 1996. Effects of clearcutting on habitat use and reproductive success of the Ovenbird in forested landscapes. *Conservation Biology* 10:1380–1386
- Kitchener DJ, Chapman A, Dell J, Muir BG, Palmer M. 1980. Lizard assemblages and reserve size and structure in the Western Australian wheatbelt Some implications for conservation. *Biological Conservation* 17:25-62
- Klingbeil BT, Willig MR. 2009. Guild-specific responses of bats to landscape composition and configuration in fragmented Amazonian rainforest. *Journal of Applied Ecology* 46:203-213

- Kumar S, Stohlgren TJ, Chong GW. 2006. Spatial heterogeneity influences native and nonnative plant species richness. *Ecology* 87:3186-3199
- Lande R. 1987. Extinction thresholds in demographic models of territorial populations. *American Naturalist* 130:624-635
- Langlois JP, Fahrig L, Merriam G, Artsob H. 2001. Landscape structure influence continental distribution of hantavirus in deer mice. *Landscape Ecology* 16:255-266
- Le Roux DS, Ikin K, Lindenmayer DB, Manning AD, Gibbons P. 2015. Single large or several small? Applying biogeographic principles to tree-level conservation and biodiversity effects. *Biological Conservation* 191:558-566
- Leopold, A. 1933. Game management. Charles Scribner Sons, New York, NY, USA
- Levins R. 1970. Extinction. pp. 77–107 in M. Gesternhaber (ed.), *Some mathematical problems in biology*. American Mathematical Society, Providence, RI, USA
- Levins R, Culver D. 1971. Regional coexistence of species and competition between rare species. *Proceedings of the National Academy of Sciences of the United States of America* 68:1246-1248
- Lichstein JW, Simons TR, Franzreb KE. 2002. Landscape effects on breeding songbird abundance in managed forests. *Ecological Applications* 12:836-857
- Lindell CA, Riffell SK, Kaiser SA, Battin AL, Smith ML, Sisk TD. 2007. Edge responses of tropical and temperate birds. *Wilson Journal of Ornithology* 119:205–220
- Lindsay KE, Kirk DA, Bergin TM, Best LB, Sifneos JC, Smith J. 2013. Farmland heterogeneity benefits birds in American Mid-west watersheds. *American Midland Naturalist* 170:121-143
- Liu W, Zhang Q, Liu G. 2012. Influences of watershed landscape composition and configuration on lake-water quality in the Yangtze River basin of China. *Hydrological Processes* 26:570-578
- Loehle C. 1987. Testing in ecology: psychological aspects and the importance of theory maturation. *Quarterly Review of Biology* 62:397-409
- MacArthur RH, Wilson EO. 1967. *The theory of island biogeography*. Princeton University Press, Princeton, NJ, USA
- Magle SB, Kardash LH, Rothrock AO. 2015. Movements and habitat interactions of white-tailed deer: implications for chronic wasting disease management. *American Midland Naturalist* 173:267-282
- Martin AE, Fahrig L. 2012. Measuring and selecting scales of effect for landscape predictors in species—habitat models. *Ecological Applications* 22:2277–2292
- Martínez-Sanz C, Canzano CSS, Fernández-Aláez M, García-Criado F. 2012. Relative contribution of small mountain ponds to regional richness of littoral macroinvertebrates and the implications for conservation. *Aquatic Conservation: Marine and Freshwater Ecosystems* 22:155-164
- Mbora DN, McPeek MA. 2009. Host density and human activities mediate increased parasite prevalence and richness in primates threatened by habitat loss and fragmentation. *Journal of Animal Ecology* 78:210-218
- McAlpine CA, Eyre TJ. 2002. Testing landscape metrics as indicators of habitat loss and fragmentation in continuous eucalypt forests (Queensland, Australia). *Landscape Ecology* 17:711-728
- McAlpine CA, Rhodes JR, Callaghan JG, Bowen ME, Lunney D, Mitchel DL, Pullar DV, Possingham HP. 2006. The importance of forest area and configuration relative to local

- habitat factors for conserving forest mammals: A case study of koalas in Queensland, Australia. *Biological Conservation* 132:153-165
- McCoy ED, Mushinsky HR. 1994. Effects of fragmentation on the richness of vertebrates in the Florida scrub habitat. *Ecology* 75:446-457
- McGarigal K, Cushman SA. 2002. Comparative evaluation of experimental approaches to the study of habitat fragmentation effects. *Ecological Applications* 12:335-345
- McGarigal K, McComb WC. 1995. Relationships between landscape structure and breeding birds in the Oregon Coast Range. *Ecological Monographs* 65:235-260
- McGarigal K. 2015. Fragstats help 4.2.
 - http://www.umass.edu/landeco/research/fragstats/documents/fragstats_documents.html
- McNeil SE, Fairweather PG. 1993. Single large or several small marine reserves? An experimental approach with seagrass fauna. *Journal of Biogeography* 20:429-440
- Meyer JS, Irwin LL, Boyce MS. 1998. Influences of habitat abundance and fragmentation on Northern Spotted Owls in western Oregon. *Wildlife Monographs* 139:3-51.
- Mitchell MS, Rutzmoser SH, Wigley TB, Loehle C, Gerwin JA, Keyser PD, Lancia RA, Perry RW, Reynolds CJ, Thill RE, Weih R, White D, Wood PB. 2006. Relationships between avian richness and landscape structure at multiple scales using multiple landscapes. *Forest Ecology and Management* 221:155-169
- Moilanen A, Nieminen M. 2002. Simple connectivity measures in spatial ecology. *Ecology* 83:1131–1145
- Moore CH, Van Neil K, Harvey ES. 2011. The effect of landscape composition and configuration on the spatial distribution of temperate demersal fish. *Ecography* 34:425-435.
- Mortelliti A, Amori G, Capizzi D, Cervone C, Fagiani S, Pollini B, Boitani L. 2011. Independent effects of habitat loss, habitat fragmentation and structural connectivity on the distribution of two arboreal rodents. *Journal of Applied Ecology* 48:153-162
- Nickerson RS. 1998. Confirmation bias: a ubiquitous phenomenon in many guises. *Review of General Psychology* 2:175-220
- Oertli B, Joye, DA, Castella E, Juge R, Cambin D, Lachavanne J-B. 2002. Does size matter? The relationship between pond area and biodiversity. *Biological Conservation* 104:57-70
- OMNR. 2002. Forest management guide for natural disturbance pattern emulation, Version 3.1. Ontario Ministry of Natural Resources, Queen's Printer for Ontario, Toronto, ON, Canada.
- Ouin A, Martin M, Burel F. 2013. Agricultural landscape connectivity for the meadow brown butterfly (*Maniola jurtina*). *Agriculture, Ecosystems and Environment* 124:193–199
- Palmer MA, Swan CM, Nelson K, Silver P, Alvestad R. 2000. Streambed landscapes: evidence that stream invertebrates respond to the type and spatial arrangement of patches. *Landscape Ecology* 15:563-576
- Peak RG, Thompson FR. 2013. Amount and type of forest cover and edge are important predictors of Golden-Cheeked Warbler density. *The Condor* 115:659-668
- Pellissier V, Cohen M, Boulay A, Clergeau P. 2012. Birds are also sensitive to landscape composition and configuration within the city centre. *Landscape and Urban Planning* 104:181–188
- Perović D, Gámez-Virués S, Bőrschig C, Klein A-M, Krauss J, Steckel J, Rothenwöhrer C, Erasmi S, Tscharntke T, Westphal C. 2015. Configurational landscape heterogeneity shapes functional community composition of grassland butterflies. *Journal of Applied Ecology* 52:505-513

- Plećaš M, Gagić V, Jancović M, Petrović-Obradović O, Kavallieratos NG, Tomanović Ž, Thies C, Tscharntke T, Ćetković A. 2014. Landscape composition and configuration influence cerealaphid–parasitoid–hyperparasitoid interactions and biological controldifferentially across years. *Agriculture, Ecosystems and Environment* 183:1–10
- Prugh LR. 2009. An evaluation of patch connectivity measures. *Ecological Applications* 19:1300–1310
- Prugh LR, Hodges KE, Sinclair ARE, Brashares JS. 2008. Effect of habitat area and isolation on fragmented animal populations. *Proceedings of the National Academy of Sciences USA* 105:20770-20775
- Quinn JF, Harrison SP. 1988. Effect of habitat fragmentation and isolation on species richness: Evidence from biogeographic patterns. *Oecologia* 75:132-140
- Quinn JF, Robinson GR. 1987. The effects of experimental subdivision on flowering plant diversity in a California annual grassland. *Journal of Ecology* 75:837-855
- Radford JQ, Bennett AF, Cheers GJ. 2005. Landscape-level thresholds of habitat cover for woodland-dependent birds. *Biological Conservation* 124:317–337
- Radford JQ, Bennett AF. 2007. The relative importance of landscape properties for woodland birds in agricultural environments. *Journal of Applied Ecology* 44:737-747
- Ranius T, Johansson V, Fahrig L. 2010. A comparison of patch connectivity measures using data on invertebrates in hollow oaks. *Ecography* 33:1–8
- Rempel RS. 2007. Selecting focal songbird species for biodiversity conservation assessment: Response to forest cover amount and configuration. *Avian Conservation and Ecology* 2:6[online]
- Richardson SJ, Clayton R, Rance BD, Broadbent H, McGlone MS, Wilmshurst JM. 2015. Small wetlands are critical for safeguarding rare and threatened plant species. *Applied Vegetation Science*: 18: 230-241
- Riffell S, Scognamillo D, Burger LW, Bucholtz S. 2010. Broad-scale relations between conservation reserve program and grassland birds: Do cover type, configuration and contract age matter? *The Open Ornithology Journal* 3:112-123
- Robertson O, Maron M, Buckley Y, McAlpine C. 2013. Incidence of competitors and landscape structure as predictors of woodland-dependent birds. *Landscape Ecology* 28:1975-1987
- Robinson GR, Quinn JF. 1988. Extinction, turnover and species diversity in an experimentally fragmented California annual grassland. *Oecologia* 76:71-82
- Robinson SK, Thompson FR, Donovan TM, Whitehead DR, Faaborg J. 1995. Regional forest fragmentation and the nesting success of migratory birds. *Science* 267:1987-1990
- Roland J. 1993. Large-scale forest fragmentation increases the duration of tent caterpillar outbreak. *Oecologia* 93:25-30
- Rosenberg KV, Lowe JD, Dhondt AA. 1999. Effects of forest fragmentation on breeding tanagers: A continental perspective. *Conservation Biology* 13:568-583
- Rüdisser J, Walde J, Tasser E, Frühauf J, Teufelbauer N, Tappeiner U. 2015. Biodiversity in cultural landscapes: influence of land use intensity on bird assemblages. *Landscape Ecology* 30:1851-1863
- Rueda M, Hawkins BA, Morales Castilla I, Vidanes RM, Ferrero M, Rodríguez MÁ. 2013. Does fragmentation increase extinction thresholds? A European-wide test with seven forest birds. *Global Ecology and Biogeography* 22:1282-1292
- Saetersdahl M. 1994. Rarity and species/area relationships of vascular plants in deciduous woods, western Norway: Applications to nature reserve selection. *Ecography* 17:23-38

- San-José M, Arroyo-Rodríguez V, Sánchez-Cordero V. 2014. Association between small rodents and forest patch and landscape structure in the fragmented Lacandona rainforest, Mexico. *Tropical Conservation Science* 7:403-422
- Sattler T, Duellil P, Obrist MK, Arlettaz R, Moretti M. 2010. Response of arthropod species richness and functional groups to urban habitat structure and management. *Landscape Ecology* 25:941-954
- Saura S, Bodin Ő, Fortin M-J. 2014. Stepping stones are crucial for species' long-distance dispersal and range expansion through habitat networks. *Journal of Applied Ecology* 51:171-182
- Silver P, Cooper JK, Palmer MA, Davis EJ. 2000. The arrangement of resources in patchy landscapes: effects on distribution, survival, and resource acquisition of chironomids. *Oecologia* 124:216-224.
- Silver P, Wooster D, Palmer MA. 2004. Chironomid responses to spatially structured, dynamic, streambed landscapes. *Journal of the North American Benthological Society* 23:69-77
- Simberloff DS, Abele LG. 1976. Island biogeography theory and conservation practice. *Science* 191:285-286
- Simberloff DS, Abele LG. 1982. Refuge design and island biogeographic theory: Effects of fragmentation. *The American Naturalist* 120:41-50
- Simberloff D, Gotelli N. 1984. Effects of insularization on plant species richness in the prairieforest ecotone. *Biological Conservation* 29:27-46
- Sitters H, Christie FJ, Stefano JD, Swan M, Penman T, Collins PC, York A. 2014. Avian responses to the diversity and configuration of fire age classes and vegetation types across a rainfall gradient. *Forest Ecology and Management* 318:13-20
- Slancarova J, Benes J, Kristynek M, Kepka P, Konvicka M. 2014. Does the surrounding landscape heterogeneity affect the butterflies of insular grassland reserves? A contrast between composition and configuration. *Journal of Insect Conservation* 18:1-12
- Smith AC, Fahrig L, Francis CM. 2011. Landscape size affects the relative importance of habitat amount, habitat fragmentation, and matrix quality on forest birds. *Ecography* 34:103-113
- Steckel J, Westphal C, Peters MK, Bellach M, Rothenwoehrer C, Erasmi S, Scherber C, Tscharntke T, Steffan-Dewenter I. 2014. Landscape composition and configuration differently affect trap-nesting bees, wasps and their antagonists. *Biological Conservation* 172:56-64
- Tabarelli M, Mantovani W, Peres CA. 1999. E€ ects of habitat fragmentation on plant guild structure in the montane Atlantic forest of southeastern Brazil. *Biological Conservation* 91:119-129
- Tewksbury JJ, Hejl SJ, Martin TE. 1998. Breeding productivity does not decline with increasing fragmentation in a western landscape. *Ecology* 79:2890-2903
- Thornton DH, Branch LC, Sunquist ME. 2011. The influence of landscape, patch, and within-patch factors on species presence and abundance: a review of focal patch studies. Landscape Ecology 26:7–18
- Tischendorf L, Bender DJ, Fahrig L. 2003. Evaluation of patch isolation metrics in mosaic landscapes for specialist vs. generalist dispersers. *Landscape Ecology* 18:41–50
- Tischendorf L, Fahrig L. 2000. How should we measure landscape connectivity? *Landscape Ecology* 15:633-641

- Trzcinski MK, Fahrig L, Merriam G. 1999. Independent effects of forest cover and fragmentation on the distribution of forest breeding birds. *Ecological Applications* 9:586-593
- Tscharntke T, Steffan-Dewenter I, Kruess A, Thies C. 2002. Contributions of small habitat fragments to conservation of insect communities of grassland-cropland landscapes. *Ecological Applications* 12:354-363
- Uuemaa R, Roosaare J, Mander Ü. 2007. Landscape metrics as indicators of river water quality at catchment scale. *Nordic Hydrology* 38:125-138
- Villard M-A, Trzcinski MK, Merriam G. 1999. Fragmentation effects on forest birds: Relative influence of woodland cover and configuration on landscape occupancy. *Conservation Biology* 13:774-783
- Villard M-A, Metzger JP. 2014. Beyond the fragmentation debate: a conceptual model to predict when habitat configuration really matters. *Journal of Applied Ecology* 51:309-318
- Virolainen KM, Suomi T, Suhonen J, Kuitunen M. 1998. Conservation of vascular plants in single large and several small mires: Species richness, rarity and taxonomic diversity. *Journal of Applied Ecology* 35:700-707
- Walter WD, VerCauteren KC, Campa H, Clark WR, Fischer JW, Hygnstrom SE, Mathews NE, Nielsen CK, Schauber EM, Van Deelen TR, Winterstein SR. 2009. Regional assessment on influence of landscape configuration and connectivity on range size of white-tailed deer. *Landscape Ecology* 24:1405-1420
- Westphal MI, Field SA, Tyre AJ, Paton D, Possingham HP. 2003. Effects of landscape pattern on bird species distribution in the Mt. Lofty Ranges, South Australia. *Landscape Ecology* 18:413-426
- With KA. 2015. Are landscapes more than the sum or their patches? *Landscape Ecology* DOI 10.1007/s10980-015-0328-8
- With KA, King AW. 1999. Dispersal success on fractal landscapes: a consequence of lacunarity thresholds. *Landscape Ecology* 14:73-82
- With KA, Pavuk DM, Worchuck JL, Oates RK, Fisher JL. 2002. Threshold effects of landscape structure on biological control in agroecosystems. *Ecological Applications* 12:52-65
- With KA, Pavuk DM. 2012. Direct versus indirect effects of habitat fragmentation on community patterns in experimental landscapes. *Oecologia* 170:517-528
- Wolff JO, Schauber EM, Edge WD. 1997. Effects of habitat loss and fragmentation on the behavior and demography of gray-tailed voles. *Conservation Biology* 11:945-956
- Xiao H, Ji W. 2007. Relating landscape characteristics to non-point source pollution in mine waste-located watersheds using geospatial techniques. *Journal of Environmental Management* 82:111-119
- Zacharias D, Brandes D. 1990. Species area-relationships and frequency Floristical data analysis of 44 isolated woods in northwestern Germany. *Vegetatio* 88:21-29
- Zharikov Y, Lank DB, Cooke F. 2007. Influence of landscape pattern on breeding distribution and success in a threatened Alcid, the marbled murrelet: Model transferability and management implications. *Journal of Applied Ecology* 44:748-759

Figure captions

- Figure 1. The concept of 'habitat patchiness' in ecology. a. From 1935 to 1970, habitat patchiness was associated with the concept of 'habitat spatial heterogeneity,' and was assumed to have a positive effect on biodiversity. b. From 1970 to present, habitat patchiness is associated with the concept of 'habitat fragmentation,' and is assumed to have a negative effect on biodiversity. This apparent contradiction is resolved by recognizing that the former is often associated with habitat gain while the latter with habitat loss.
- Figure 2. Habitat loss is a process whereby habitat is destroyed over time. In contrast, habitat fragmentation *per se* (hereafter 'habitat fragmentation') is a difference in spatial pattern. For a given amount of habitat, a more fragmented pattern has more, smaller patches, with more total edge in the landscape. The current dominant paradigm assumes that habitat fragmentation generally has negative effects on biodiversity. If this is true, then policies should favour 'land sparing' over 'land sharing.'
- Figure 3. Patch isolation is an inverse measure of habitat amount. Patch A is less isolated than Patch B precisely because there is more habitat in the local landscape around Patch A than in the local landscape around Patch B.
- Figure 4. Illustration of the difference between a patch-scale study and a landscape-scale study. a. In a patch-scale study, predictor variables are spatial attributes of individual patches, e.g. patch size, patch isolation. b. In a landscape-scale study, predictor variables are spatial attributes of individual landscapes, e.g. percentage of the landscape in habitat, habitat fragmentation.
- Figure 5. a. Experimental study design for evaluating the independent effects of habitat amount and habitat fragmentation. b. Aerial photograph of a landscape-scale experiment where landscapes were constructed such that aphid habitat fragmentation was varied (2 levels) independent of habitat amount (6 levels), with 3 replicates of each combination of habitat fragmentation and habitat amount. Habitat is planted clover, and each landscape is 16 m x 16 m (from With et al. 2002).
- Figure 6. SLOSS-type study. a. Species are sampled in each patch within a global set of patches. b-d. Subsets of the global set, such that each subset has the same total habitat amount for all patches combined, but subsets vary in habitat fragmentation.
- Figure 7. SLOSS-type analysis. Species number is accumulated across patches, from the largest to smallest patch (grey points and lines) and from the smallest to largest patch (black points and lines). When the largest-to-smallest curve is above the smallest-to-largest curve (A), fragmentation has a negative effect on species richness. When the reverse is true (B), fragmentation has a positive effect on species richness. When the lines are co-incident or cross each other (C), there is no clear effect of fragmentation on species richness.
- Figure 8. Proportion of significant fragmentation effects that are negative and positive, across all ecological responses. Numbers above bars are the numbers of significant effects. Most significant fragmentation effects are positive.

Figure 9. Proportion of significant fragmentation effects that are negative and positive. Numbers above bars are the numbers of significant effects. Significant effects of habitat fragmentation are mostly positive, irrespective of (a) the method used to control for habitat amount effects, (b) the response taxon, (c) the fragmentation metric used, (d) the category of response variable, and (e) species IUCN status. (a) For a description of the three methods for controlling for the effects of habitat amount see section 2, How to measure effects of habitat fragmentation independent of habitat amount. (b) Six significant responses to fragmentation combined two or more taxonomic categories; these were excluded here. 'Water quality' included variables such as abundances of water quality indicator taxa and water chemistry variables such as nutrient concentration in receiving waters; a significant negative effect of fragmentation on nutrient concentration was interpreted as a significant positive effect on water quality. (c) Infrequently used fragmentation metrics were re-labeled according to the common metric to which they were most strongly related. For example, 'splitting index' was re-labeled as 'number of patches', 'like adjacencies' as 'clumping', and 'mean circumscribing circle' as 'edge density'. When multi-variate methods were used to create landscape measures, the composite variable (e.g. PCA axis) representing fragmentation was re-labeled as the metric most strongly correlated to it. For SLOSS-type analyses the metric was 'number of patches'. 'Whole landscape metrics' were those measured over all cover classes, i.e. 'landscape metrics' in McGarigal (2015). A significant positive response to a 'clumping' metric was recorded as a significant negative fragmentation effect (and vice-versa). (d) 'Multispecies' includes mainly species richness but also species diversity measures and total cross-species abundance. 'Multispecies - specialist/threatened' responses are a subset of 'multispecies' significant responses, where the species were limited by the authors to threatened and/or specialist species. 'Single species - movement' and 'single species demographic' responses are subsets of 'single species' significant responses; most of the other 'single species' responses were species abundance or occurrence. (e) IUCN status (IUCN 2015) includes single species responses only. "Threatened" includes 2 species listed as endangered, one showing a negative and one showing a positive significant fragmentation response, and 5 near near-threatened species all showing significant positive fragmentation responses. 'declining' / 'stable' / 'increasing' are estimated species population trends.

Figure 10. The effects of habitat clumping and habitat amount are confounded when the species response combines data from multiple sample sites (black dots) within habitat, in each landscape. In landscapes containing more clumped habitat (b), a larger proportion of sample sites are in parts of the landscape containing more habitat in their 'local landscapes' (circles surrounding sample sites) than in landscapes containing less clumped habitat (a).

Figure 11. Proportion of significant fragmentation effects that are negative and positive. Numbers above bars are the numbers of significant effects. No support was found for any of three hypotheses predicting situations in which the effects of habitat fragmentation should be strongly negative. (a) Significant responses were not more likely to be negative in tropical systems. (b) Larger-ranging species or species groups were not more likely to show significant negative responses to fragmentation. Here, I indexed movement range as the maximum radius (in m) of the individual replicate landscapes considered by the authors, on the assumption that authors chose to evaluate the effects of fragmentation over larger spatial extents for species with larger movement ranges. (c) Significant effects of habitat fragmentation were not more likely to

Fahrig L. 2017 (in press). Annual Reviews of Ecology, Evolution and Systematics 48.

be negative at levels of habitat amount lower than 20-30% of the landscape. Habitat amount was indexed as the median habitat amount across landscapes in each study, where 'median' was the actual median, or the mean when the median was not given, or the midpoint in the range when neither the median nor the mean was given. Note median habitat amount could not be determined in situations where landscape area increased with fragmentation (e.g. SLOSS-type analyses), or when fragmentation was a whole-landscape metric; these responses were excluded here.

Figure 12. Explanations offered by the authors, for significant positive effects of fragmentation. Numbers above bars are the numbers of significant effects. For details, see section 5. Authors' explanations for significant positive effects of habitat fragmentation.

Figure 13. Illustration of how habitat fragmentation increases landscape complementation. Landscape complementation is higher when different required resources are more accessible to each other in the landscape. Habitat fragmentation increases interspersion of different resources in the landscape, which increases their accessibility from each other (black arrows).

Figure 1.

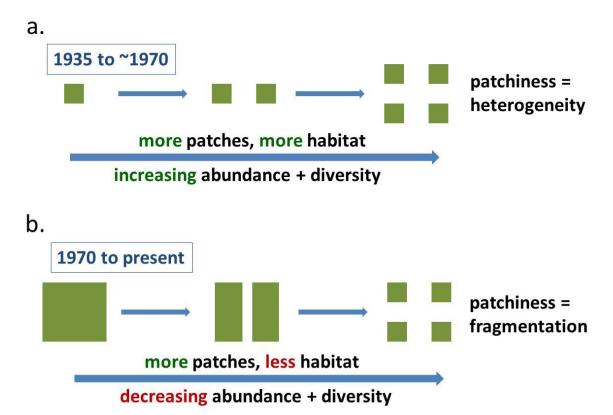


Figure 2.

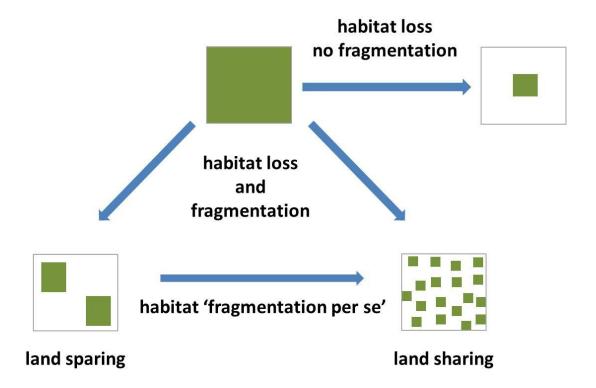


Figure 3.

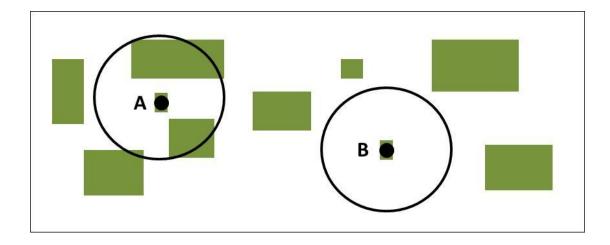
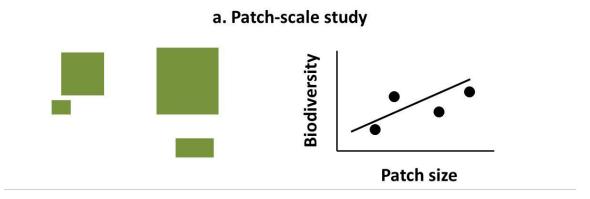


Figure 4.



b. Landscape-scale study

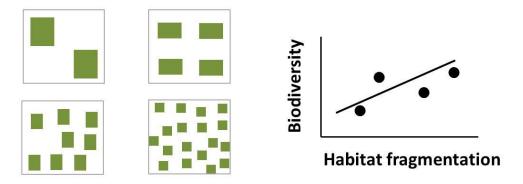


Figure 5.

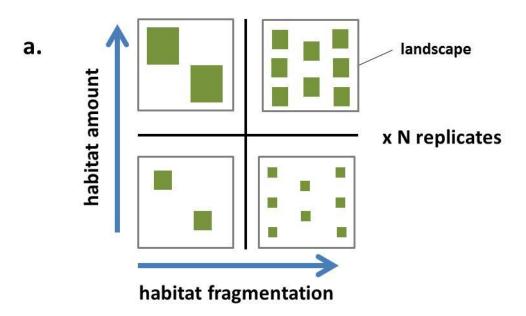
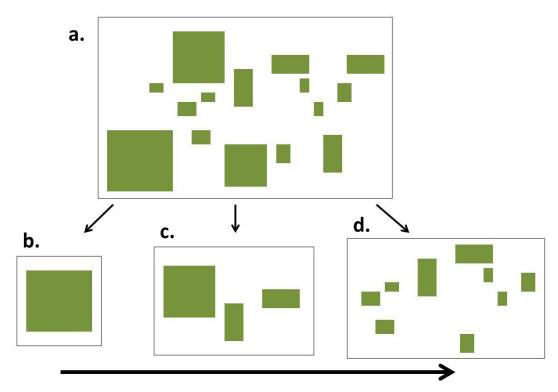


Figure 6.



constant habitat amount, increasing fragmentation per se

Figure 7.

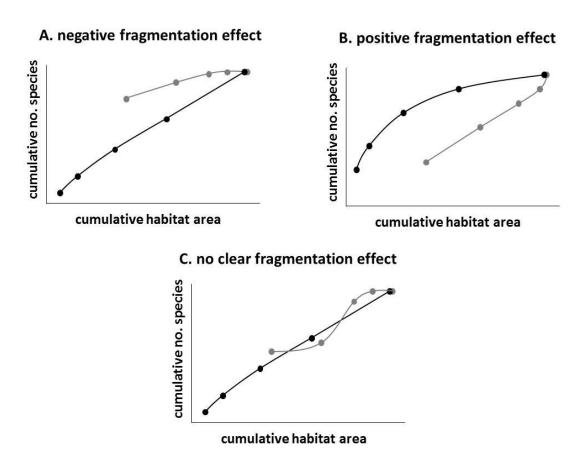


Figure 8.

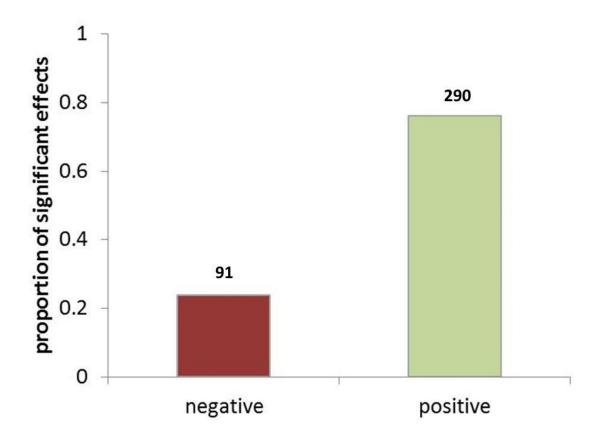


Figure 9.

a. method used to control for habitat amount effects

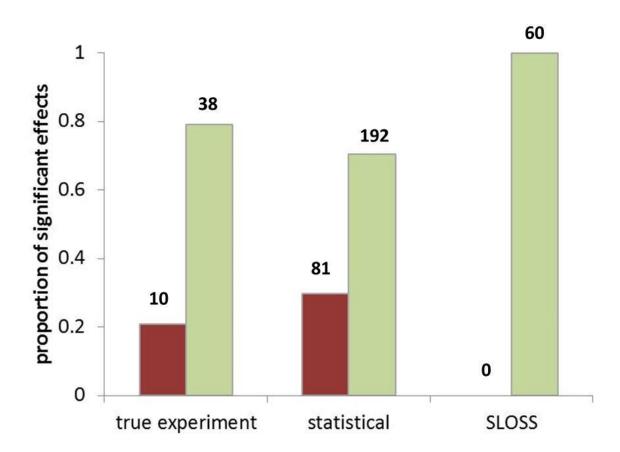


Figure 9 cont'd

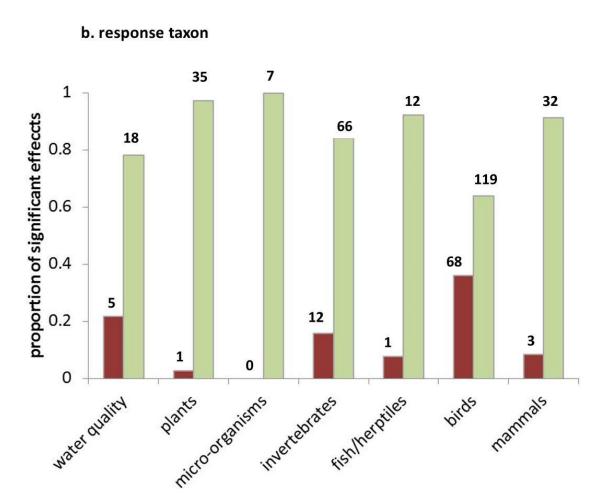


Figure 9 cont'd

c. category of fragmentation metric used

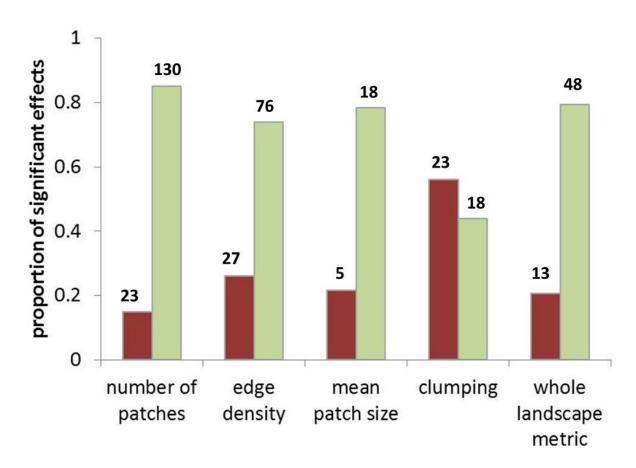


Figure 9 cont'd

d. category of response variable

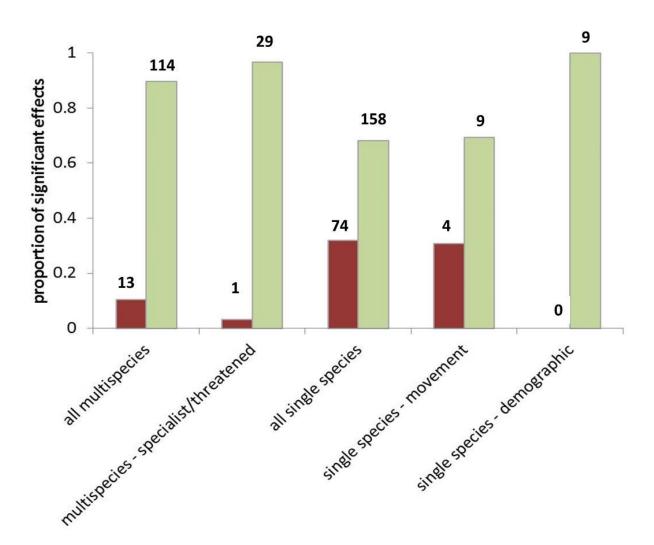


Figure 9 cont'd

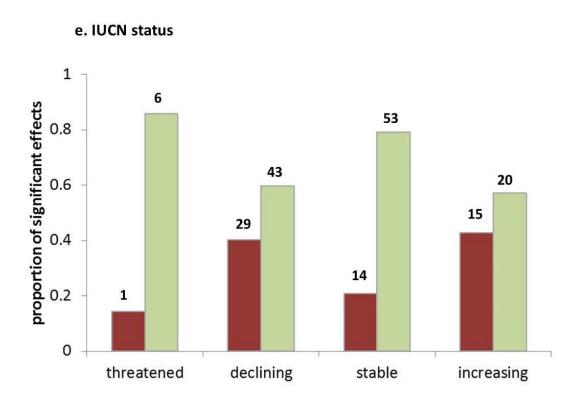
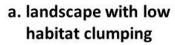
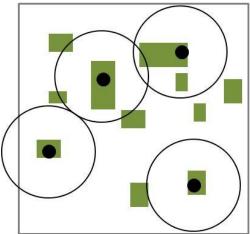


Figure 10.





b. landscape with high habitat clumping

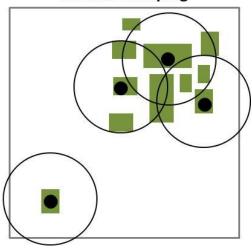


Figure 11.

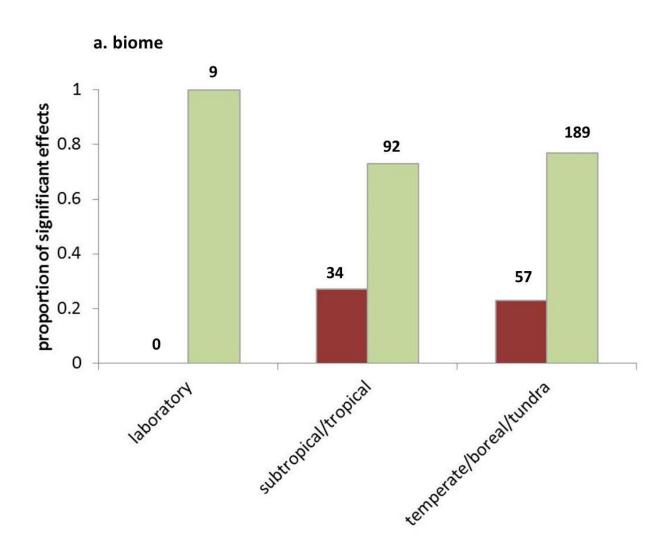


Figure 11 cont'd.

b. replicate landscape size

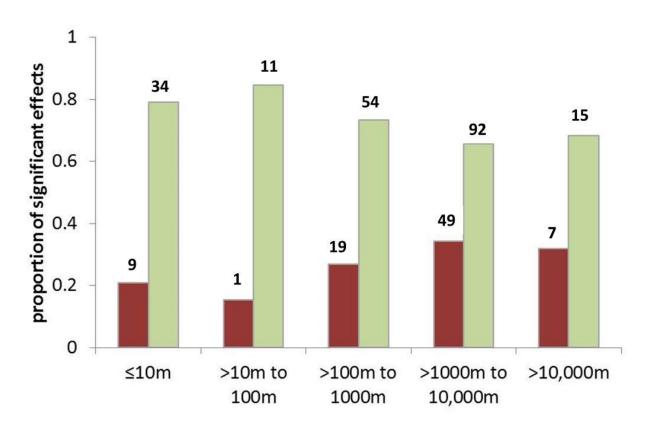


Figure 11 cont'd.

c. median habitat amount

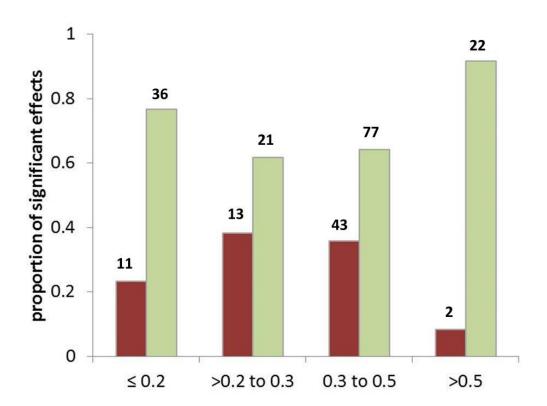


Figure 12.

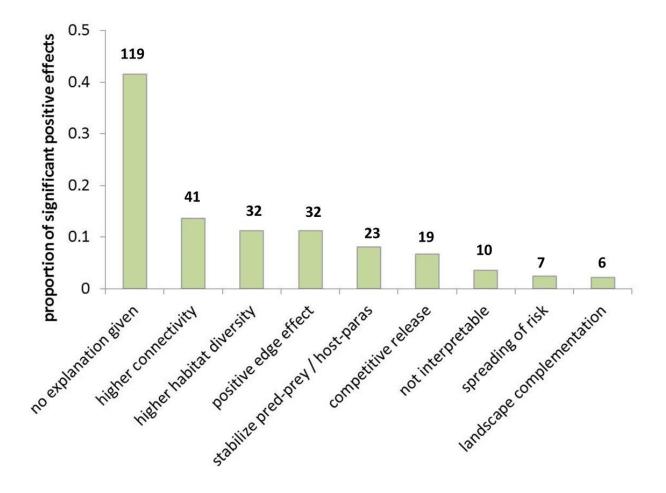


Figure 13.

