

1 **Into the wild: microbiome transplant** 2 **studies need broader ecological reality**

3 4 Authors

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26 27 **Keywords**

28 conservation, ecological adaptation, ecophysiology, holobiont, metacommunity,
29 ecosystem on a leash

37 **Abstract**

38

39 Gut microbial communities (microbiomes) profoundly shape the ecology and
40 evolution of multicellular life. Interactions between host and microbiome appear to be
41 reciprocal, and ecological theory is now being applied to better understand how hosts
42 and their microbiome influence each other. However, some ecological processes that
43 underlie reciprocal host-microbiome interactions may be obscured by the current
44 convention of highly-controlled transplantation experiments. Although these approaches
45 have yielded invaluable insights, there is a need for a broader array of approaches to
46 fully understand host-microbiome reciprocity. Using a directed review, we surveyed the
47 breadth of ecological reality in the current literature on gut microbiome transplants with
48 non-human recipients. For 55 studies, we categorized 9 key experimental conditions
49 that impact the ecological reality (EcoReality) of the transplant, including host taxon
50 match and donor environment. Using these categories, we rated the EcoReality of each
51 transplant. Encouragingly, the breadth of EcoReality has increased over time, but some
52 components of EcoReality are still relatively unexplored, including recipient host
53 environment and microbiome state. The conceptual framework we develop here maps
54 the landscape of possible EcoReality to highlight where fundamental ecological
55 processes can be considered in future transplant experiments.

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57 **A Quest for Ecological Reality**

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***We shall not cease from exploration
And the end of all our exploring
Will be to arrive where we started
And know the place for the first time.***

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T.S. Eliot - Little Gidding (1)

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65 Far from passive passengers, resident microbial communities (microbiomes) are
66 integral to the basic biological functioning of multicellular life. This revelation, ushered in
67 by advances in sequencing and computing technology, is grounded in a growing
68 understanding that microbiomes profoundly shape their host's biology, influencing
69 factors such as immunity (2), adiposity (3), thermogenesis (4), hormonal regulation (5),
70 physiological development (6), memory (7), and behaviour (8). To date, highly-
71 controlled experiments with laboratory rodent microbiomes have provided foundational
72 and indispensable knowledge on host-microbiome interactions. Furthermore, these
73 initial experiments have set the stage for integrative contributions by comparative
74 animal physiologists, ecologists, and evolutionary biologists to fill knowledge gaps in our
75 understanding of host-microbiome evolution and the interactions which underlay these

76 partnerships (9).

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78 Recently, researchers have started to appreciate the intertwining nature of host-
79 microbiome interactions. Evidence is mounting that hosts can shape the composition of
80 their microbiome community (10), and that microbiomes can influence their host's
81 behaviour (8) and physiology (5). Based on differing cases of how host and microbiome
82 might interact, Foster et al. (11) proposed four distinct models: 1) 'host control', in which
83 the host unilaterally governs the composition of its microbiome; 2) 'symbiont control', in
84 which the microbiome shapes the host phenotype; 3) 'open ecosystem', in which the
85 host and microbiome do not interact; and 4) 'ecosystem on a leash', in which the host
86 influences the microbiome by selecting upon microbial function rather than for specific
87 microbial taxa. These connections can be so intimate that some researchers (12,13)
88 proposed that a host and its associated microorganisms are a single biological entity—
89 termed the 'holobiont'—on which selection acts, challenging notions of organismal
90 individuality. Using this holobiont perspective, Alberdi et al. (14) posited that the
91 microbial component of the holobiont, with its greater mutability compared to the host
92 genome, may be an important mechanism facilitating host adaptation to rapid
93 environmental change. Therefore, understanding the interplay between the host and the
94 microbiome is crucial for addressing both fundamental and applied questions about the
95 microbiome.

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97 Host-microbiome interactions are shaped by ecological and evolutionary
98 processes (15,16). Because host-microbiome interactions are potentially reciprocal,
99 these processes act on three levels: the assembly and dynamics of the microbiome, the
100 influence of the host on the microbiome, and the influence of the microbiome on the
101 host. Microbiome assembly is governed by a variety of factors including environmental
102 filtering, priority effects, random sampling, and dispersal limitation (16,17). The within-
103 microbiome community dynamics are influenced by new invasions, competition,
104 mutualisms, and other interactions (15). A host's actions can also shape their
105 associated microbiomes. For example, the host's social behaviour can impact microbial
106 dispersal (18). Conversely, the dynamics of the microbiome can impact the host; the
107 change in microbiome community composition leading to *Clostridium difficile*
108 colonization and pathogenicity is a classic example in humans (19). Evolutionary
109 processes also occur in tandem with all the ecological processes mentioned previously
110 because of the short timescales associated with microbial turnover relative to microbial
111 evolutionary rates (17). Consequently, considering the ecological processes that
112 underlie host-microbiome interactions is critical for making sense of the reciprocity
113 between the host and its microbiome.

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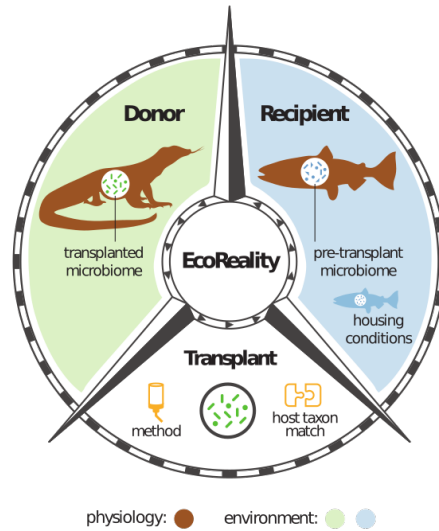
115 The most convincing evidence for host-microbiome interactions has been

116 gleaned through microbiome transplantation studies. In these studies, researchers
117 experimentally translocate microbial species or communities from donor hosts or
118 external substrates to recipient hosts. Highly-controlled transplantation studies have
119 been and will continue to be invaluable to experimentally probe the host-microbiome
120 relationship. However, there is a trade-off: highly-controlled experiments isolate
121 mechanisms of interest, but they cannot simultaneously capture the full suite of
122 ecological processes (drift, dispersal, competition, etc.) that influence reciprocal host-
123 microbiome interactions in nature. For example, the use of germ-free recipients may
124 preclude competition between introduced and resident microbes (20), and isolated
125 laboratory conditions may limit the potential for microbial dispersal from influencing the
126 composition of the resulting microbiome (21). How researchers weigh this trade-off
127 depends on the research question of interest. If a researcher's goal is to understand the
128 effect of specific microbes on host physiology, or to develop applications for human
129 health and domestic animal production, controlled experimental conditions are
130 preferred. In contrast, when examining the role and consequences of reciprocal host-
131 microbiome interactions in ecological and evolutionary contexts (e.g., fitness effects,
132 intergenerational microbial transmission, speciation, species persistence, etc.),
133 ecological complexity needs to be considered (22). Therefore, a comparison of highly-
134 controlled transplants and ecologically realistic (which we term EcoReal, see Box 1 for a
135 full definition) transplants that match what the host plus its microbiome would
136 experience in a wild ecosystem is required. The trade-offs of laboratory approaches and
137 the need for comparison to studies that use ecologically realistic conditions have long
138 been recognized by comparative animal physiologists (23), though to date there does
139 not seem to have been a similar recognition in microbiome research. Specifically, the
140 breadth of EcoReality in microbiome transplant studies has not been examined,
141 meaning such an evaluation remains an exciting potential avenue for future work.

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143 Here, we probe the current EcoReality of microbiome transplantation studies.
144 Our work here is not unlike Hanage's (24) questioning of the reality and applicability of
145 biomedical microbiome studies. By taking advantage of the recent explosion of studies
146 conducting microbiome transplants, we evaluated whether the current microbiome
147 transplant literature limits opportunities for ecological processes to influence study
148 outcomes. We investigated two key questions: 1) how EcoReal are the experimental
149 conditions in the current microbiome transplant literature? and 2) does the literature
150 currently cover the full potential range of EcoReality? Using long-established ecological
151 concepts, we categorized microbiome transplantations into different experimental
152 conditions which can impact the EcoReality of the transplant (Figure 1 and Box 2).
153 Using this framework, we scored the EcoReality of microbiome transplant studies that
154 used non-human recipients. We show that, overall, the breadth of EcoReality of the
155 present microbiome transplant literature has increased over time. However, EcoReality

156 has been constrained by hosts bred and kept in lab conditions, and with transplants into
 157 germ-free recipient hosts. Importantly, we provide a conceptual framework, illustrated in
 158 Figure 1, to help broaden the range of EcoReality in transplant experiments and to
 159 facilitate comparisons between transplants of varying EcoReality.

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176 Figure 1: Conceptual framework of all the experimental conditions in a microbial
 177 transplant where EcoReality can vary. See Box 2 for explanations for each experimental
 178 condition.

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180 **Box 1. Key terms and definitions**

Term	Definition
Transplant Instance	A transplant of a microbial strain or community from its native host or substrate to a different host population. A given study can involve multiple transplant instances, which are delineated based on non-substitutability of host populations or of transplant parameters.
Experimental Conditions	A decision or step in a transplant instance where there is the potential for variation in ecological reality.
Level of EcoReality	The degree to which an experimental condition matches the conditions that a host-microbiome

interaction would experience in a wild ecosystem. Each experimental condition possesses its own intrinsic EcoReality. Each transplant instance can also be assigned an EcoReality score.

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182 **Box 2.** Ecological reasoning for each experimental conditions within a transplant

Experimental condition	Reasoning Ecological theory + application to microbiome
Taxon Match	Organisms can become locally adapted (25). Local adaptation of a microbial species to a given host may mean it is not adapted to hetero-specific hosts and will perform poorly after transplantation (26).
Donor & Recipient Environment	During community assembly, the local environment acts as a filter, incorporating species from a wider species pool (21). From the microbiome’s perspective, the host’s physiology and the external environment are one intertwined environment. Therefore, the external environment can affect microbiome dynamics in two ways: indirectly through impacting the host physiology (27), and directly through the wider microbial species pool that the host and its microbiome has access to.
Donor & Recipient Physiology	The local environment acts as a filter in community assembly (21). For this experimental condition, we define physiology as physiological states that would occur regardless of the external environmental context (e.g. gene knock-out, disease-state). We differentiate intrinsic physiology from mutable host physiological responses to the external environmental context. Although these indirect environmental effects acting through host physiology are relevant (27), they are captured by the ‘Environment’ experimental condition. A host’s physiology is the <i>de facto</i> environment of inhabitant microbes, and changes or dysregulation in the host may disrupt associations between host and the microbes that persist under homeostatic physiological conditions.
Transplanted Microbiome	The interactions within an invading community, including predation or mutualism, can impact whether colonisation is successful or not (20). Thus, a full community microbiome

	transplantation may differ significantly from the transplantation of a single microbe monoculture at artificially high densities.
Transplant Method	Species have different dispersal abilities (27) and local environments filter species from the wider species pool (28). Active transplantations may circumvent differing dispersal abilities of microbial species and may undermine host filtering of the microbial community. Furthermore, active transplant methods can stress the host thereby changing host physiology and disrupting endogenous microbial communities (29).
Recipient Pre-transplant Microbiome	High species diversity in a community is predicted to reduce niche opportunities and to increase invasion resistance (20). Germ-free or antibiotic perturbed recipients are likely to have lower invasion resistance than recipients with intact microbiomes.
Housing conditions	Dispersal between patches is an integral ecological process which can maintain stable populations or can rescue extirpated populations (21,30). Recipient host cohabitation allows for further transmissions of the microbiome.

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184 **Lay of the land**

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186 *Literature Search*

187 We conducted a directed review of the existing literature on gut microbiome
188 transplants, finishing on October 26th 2018. We conducted our literature search in three
189 stages. First, to gauge the extent of the current literature, we did a preliminary search of
190 gut microbiome transplant studies using both Google Scholar and Web of Science
191 (University of Guelph subscription). Based on this preliminary search, we conducted a
192 more methodical search using both Google Scholar and Web of Science. Search terms
193 can be found in the Supporting Information (SI). We then sought additional publications
194 through searching the citations of papers already collected using the Web of Science
195 citations tool. We retained only those studies that conducted at least one gut
196 microbiome transplant into a non-human recipient organism. To ensure our findings
197 were generalizable to ecological and evolutionary frameworks across a broad range of
198 taxa and ecosystems, we excluded studies focused on a single human disease, *C.*
199 *difficile*.

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202 *Literature Evaluation*

203 For each study that met our criteria, we determined the number of transplant
204 instances, which we defined as the transfer of a microbial strain or community from its
205 native host or substrate to a different host population (see Box 1). We used transplant
206 instances as our unit of focus because many studies contained multiple transplant
207 instances which sometimes differed substantially in EcoReality (e.g. Seedorf et al. (31)).
208 For studies that had sequential transplants (i.e., transplant from donor to a first
209 recipient, which then was the donor for a second recipient, e.g. Seedorf et al. (31)), we
210 used only the first phase of the transplant experiment.

211 We identified nine key experimental conditions in a transplant where variation in
212 EcoReality might substantially affect the outcome of the experiment: host taxon match,
213 donor environment, donor physiology, transplanted microbiome, transplant method,
214 recipient pre-transplant microbiome, recipient environment, recipient physiology, and
215 recipient housing conditions (see Figure 1 & Box 2). Each experimental condition was
216 given an ordinal data scale (see SI Table S1) based on the range of observed and
217 possible levels for that condition, with one always representing the lowest level of
218 EcoReality. Our goal was to maintain similar resolution for each highly dimensional
219 experimental condition within our framework. For example, Taxon Match could have
220 included more levels to capture phylogenetic distance, geographic distance, and
221 feeding relationships between host and recipient (32). However given the variation in
222 scale, generalization across host taxa would have then been difficult. The levels in each
223 experimental condition were based on likely conditions found in the wild. For example,
224 with respect to the Transplanted Microbiome experimental condition, a single bacterial
225 strain at high densities entering a host in the wild is less likely than invasion by mixed
226 communities. For each transplant instance, we characterized the level of EcoReality in
227 each of the 9 experimental conditions. To ensure consistent evaluation methods,
228 EcoReality scores for each transplant instance were determined independently by two
229 co-authors (separate pairs of co-authors randomized per paper). The co-author pairs
230 then compared their scores and agreed upon the final transplant EcoReality scores.

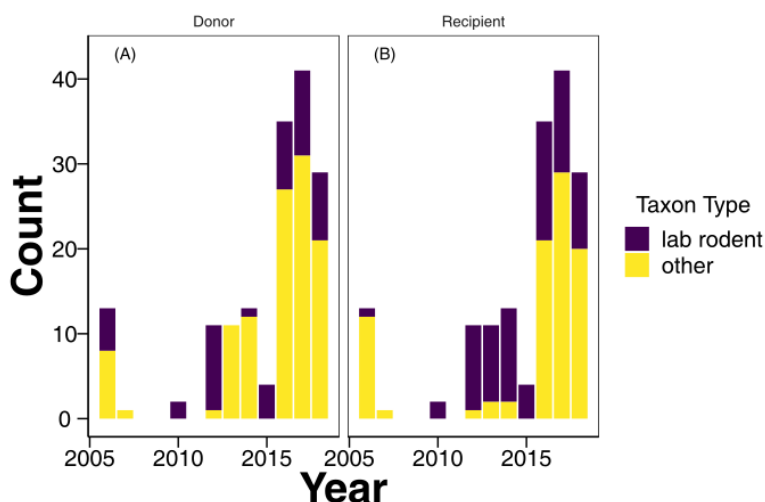
231 To determine the overall standardized EcoReality score of a transplant instance,
232 we divided each score by its corresponding maximum potential EcoReality score and
233 then added the scaled scores for each experimental condition. Thus all experimental
234 conditions were equally weighted in the overall calculation of standardized EcoReality.

235 We separated lab rodents from other animals in our results for each experimental
236 condition because the ecology, physiology and genetics of lab-reared, inbred rodent
237 models are heavily modified from wild-type rodents and other wild animals in ways that
238 may affect our understanding of reciprocal host-microbiome interactions (for example
239 Newman et al. (23) and Walter et al. (33)).

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242 *Literature EcoReality patterns*

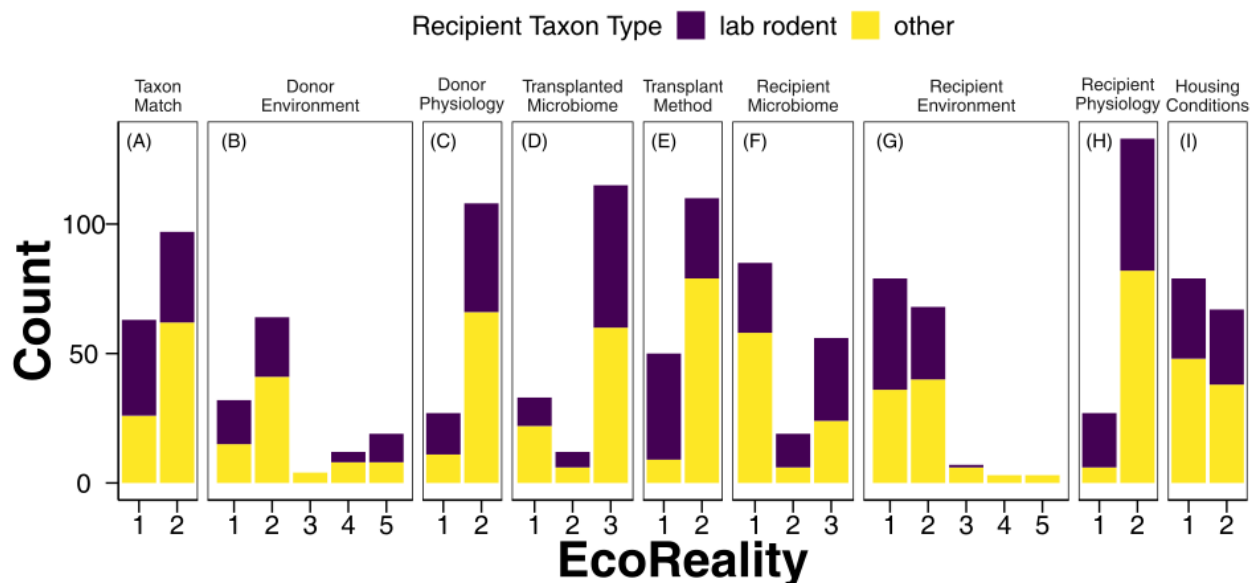
243 Our literature search returned 55 articles that met our criteria for inclusion. These
244 articles ranged from having one to 13 transplant instances with an average of 2.91
245 transplant instances per article and a total of 160 from all articles. There was a clear
246 shift over time in the number of articles using microbiome transplants. Notably, there
247 were 20 articles in the first 10 years of our search period in comparison to almost 40
248 articles during 2015-2018 (SI Figure 1). This increase coincided with a switch from
249 mainly lab rodent studies to a more diverse group of donor hosts (Figure 2A, around
250 2013), and later also to more diverse recipient hosts (Figure 2B, around 2016).



266 Figure 2: Number of transplant instances over time where the donor or recipient animal
267 was either a lab rodent (mouse or rat) or another animal.

268
269 The transplant conditions Donor and Recipient Physiology had the highest
270 EcoReality with average scores of 1.8 out of 2 (Figure 3C & H). Taxon match (score 1.6
271 out of 2, Figure 3A), transplanted microbiome (score 2.5 out of 3, Figure 3D), transplant
272 method (score 1.7 out of 2, Figure 3E), and housing condition (score 1.5 out of 2, Figure
273 3I) were moderately EcoReal. Donor environment (score 2.4 out of 5, Figure 3B),
274 recipient environment (score 1.6 out of 5, Figure 3G), and recipient microbiome (score
275 1.8 out of 3, Figure 3F) had the lowest EcoReality. Breaking EcoReality into recipient
276 lab rodents and other animals, we see that active transplant methods (score of 1) were
277 used more for lab rodents, and passive transplant methods (score of 2) were used more
278 for other animals (Figure 3E). Interestingly, there were fewer transplants with germ-free
279 recipient lab rodents than germ-free recipient other animals (score of 1) (Figure 3F).
280 This pattern was driven by bees (19 out of 85 transplant instances from five articles)
281 and zebrafish (14 out of 85 transplant instances from two articles). Overall, most
282 transplants were performed with matching (score of 2, Figure 3A) wild-type, non-

283 diseased donor and recipient hosts (score of 2, Figures 3C & H) using passive
 284 transplant methods (score of 2, Figure 3E) of whole microbial communities (score of 3,
 285 Figure 3D) and with a mixture of individual (score of 1, Figure 3I) and cohousing (score
 286 of 2, Figure 3I) of recipient hosts. However, transplants were mostly in sterile or normal
 287 lab conditions (score of 1 & 2, Figures 3B & G) with germ-free recipient hosts (score of
 288 1, Figure 3F).
 289



290 Figure 3: Number of transplant instances in each experimental condition, separated into
 291 whether the recipient animal was a lab rodent or another animal. The X-axis is the level
 292 of EcoReality, with 1 always the lowest EcoReality. The levels are explained in Table S1
 293 of our Supporting Information.

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295 Although the maximal EcoReality score increased, increasing the breadth of
 296 EcoReality studied, the maximal EcoReality score was still below the theoretical
 297 maximum standardized EcoReality score of 9 possible outlined in our framework (Figure
 298 4).

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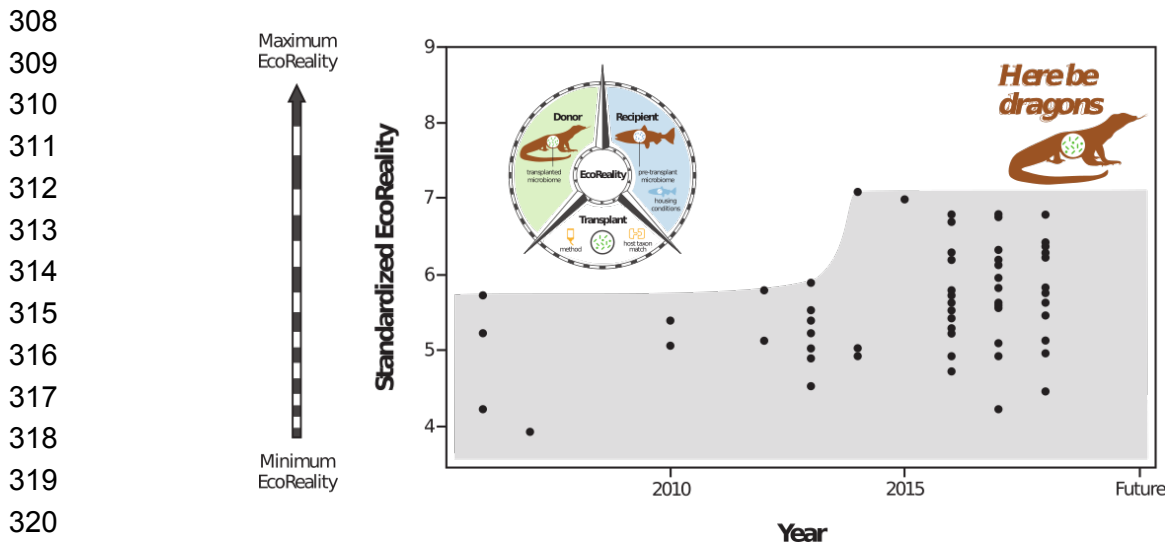


Figure 4: Standardized EcoReality score for each transplant instance. The grey area identifies the zone of EcoReality that has been studied in the literature, and the “Here be Dragons” area is the unexplored zone of EcoReality that is bound at the top by the theoretical maximum standardized EcoReality score of 9.

Here Be Dragons!

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The burgeoning field of microbiome research is integrating the traditionally disparate disciplines of ecology, evolution, and physiology, which examine distinct but interrelated processes at different scales. Yet, these interrelated processes across scales are inherent in host-microbiome relationships (e.g. Stothart et al. (34)), and thus further integration of ecology, evolution, and physiology with microbiology will be crucial for unlocking important insights about the interactions between hosts and their microbiome. As microbiome research expands further to include ecological processes that are well established in traditional ecosystems, studies that can capture these processes will be necessary. Here, we expand on the insights from foundational highly-controlled experiments that identified key mechanisms in host-microbiome interactions. We surveyed the state of the microbiome transplant literature and identified gaps in how well ecological processes are captured in transplants, what we term as ecological reality, (i.e., EcoReality, see Box 1). Our results are promising; the breadth of EcoReality is increasing over time in transplant experiments (Figure 4), but there are still some key gaps in the types of studies conducted on host-microbiome interactions (Figure 3). We suggest that a critical step in understanding reciprocal host-microbiome interactions includes explicitly designing a broader array of studies that can evaluate the role of various ecological processes that are known to shape traditional ecological systems.

348

349 Our evaluation of the microbiome transplantation literature revealed broadening
350 EcoReality in experimental procedures. Lately, there has been a sharp increase in
351 taxonomic diversity of both donor and recipient hosts (Figure 2). Transplants often used
352 passive transplantation methods with wild-type non-diseased donors as well as a
353 mixture of individual and cohabitation housing conditions (Figure 3). Finally, the
354 maximal EcoReality score of microbiome transplant studies has increased over time
355 (Figure 4). These results are encouraging because they suggest that researchers are
356 building on the initial flurry of highly-controlled transplant experiments and designing
357 diverse studies that differ in their degree of EcoReality in several of the categories we
358 examined. Continuing to broaden EcoReality will be essential for understanding the
359 ecological and evolutionary processes at work in reciprocal host-microbiome
360 interactions.

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362 However, our results show that the current literature lacks EcoReality in two key
363 areas: host environment and the state of the recipient microbiome (Figure 3). Although
364 the environment of the donor hosts was on average more EcoReal than the
365 environment of the recipient hosts, the EcoReality of the donor and recipient host's
366 environments was generally low. Most studies that we evaluated used laboratory
367 settings that exclude the chance for hosts to encounter the broader microbial species
368 pool in the environment (18,21). Laboratory conditions can also either increase or
369 decrease conspecific interactions relative to what would be observed in nature, thus
370 affecting the dispersal of microbes between hosts (18). Furthermore, laboratory
371 conditions may be obscuring feedbacks between the host and its microbiome that can
372 impact diet and habitat choice (18). The second key area lacking EcoReality is the state
373 of the recipient microbiome where most recipient hosts were germ-free. Although some
374 animals naturally start out with germ-free gastrointestinal tracts (e.g., newly eclosed
375 worker bees (35)) or do not have a resident microbiome (36), most animal species host
376 substantial microbial communities (37). Germ-free gastrointestinal tracts may lack key
377 biotic processes such as predation, competition, and facilitation, which are important
378 filters in classic ecological communities that act to mediate incoming species (20,38).
379 Overall, neglecting natural environments and intact recipient microbiomes risks
380 constraining the fundamental processes that impact reciprocal host-microbiome
381 interactions.

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383 Consequently, we advocate for more breadth in EcoReality in microbiome
384 transplant experiments. This breadth includes highly-controlled laboratory transplants,
385 which offer critical points of comparison, and provide a focused understanding of
386 particular mechanisms. The wider breadth of EcoReality for which we are advocating for
387 requires that we venture into the largely untested realm of highly EcoReal experimental

388 conditions (Figure 4), despite the logistical challenges likely associated with wild
389 conditions. There are many ways in which we might venture beyond our present
390 frontier. For example, using wild-caught animals that are either allowed to roam freely or
391 are housed in outdoor enclosures, and to use recipient animals with intact microbiomes,
392 rather than germ-free microbiomes (Figure 3). We also suggest identifying and
393 addressing the major phylogenetic gaps in the tree of life for the donor and recipient
394 host taxa. Overall, we call for a balance of studies dealing with all permutations of
395 EcoReality in each experimental condition. We hope researchers will use and adapt our
396 conceptual framework (Figure 1) in their own systems to incorporate EcoReality and,
397 where appropriate, consider how constrained EcoReality may impact their conclusions.
398 Likewise, we encourage researchers to report the methodological details pertaining to
399 each experimental condition we have identified. We hope that our literature evaluation
400 and conceptual framework will stimulate new avenues of collaborative research that will
401 evaluate the role of ecological processes in host-microbiome interactions.
402

403 Our literature evaluation suggests that we may understand only a small subset of
404 possible reciprocal host-microbiome interactions impacting our ability to assess the
405 conservation potential of the microbiome. Because we are presently likely constraining
406 fundamental ecological and evolutionary processes, host-microbiome studies may be
407 biased towards results that indicate a strong role of the microbiome on the host. Yet
408 researchers have already made strong and general assertions about the role of the
409 microbiome in the biology of the host. Due to the large effects of the microbiome on its
410 host and its mutability, Alberdi et al. (14) argued that the microbiome could act as an
411 additional axis of ecological adaptation for hosts. If the microbiome does act as an
412 additional axis, conserving microbial diversity and using bioaugmentation tools
413 (probiotic therapy and transplantation of microbiomes) would then be critical tools for
414 animal conservation (39,40). We caution that experimental protocols that lack
415 EcoReality might lead us to overestimate the capacity for microbiome variation to shape
416 host phenotypes in nature by biasing our understanding of the host-microbiome
417 relationship towards models of symbiont control (11). We suspect that a full reckoning of
418 the spectrum of EcoReality in microbiome transplant studies will uncover more
419 examples of the 'ecosystem on a leash' model (11), which posits an important but more
420 limited reciprocity between the host and the 'ecosystem' of the microbiome. These sorts
421 of nuanced interactions may or may not include the large microbiome effects which
422 underpin the ecological adaptation and conservation arguments above. Thus, we may
423 not yet have the level of understanding about reciprocal host-microbiome interactions
424 that is required to know the role of the microbiome in host adaptation or to confidently
425 inform conservation efforts. Moving forward, we assert that a consideration of
426 EcoReality is required in the design and interpretation of every study that explores how
427 the host-microbiome relationship impacts ecological adaptation.

428

429 Microbiome research has undoubtedly fascinated biologists across disciplines,
430 prompting advances in both pure and applied research and raising questions about
431 some of the most fundamental ideas in biology (13). Yet, the lay of the land in terms of
432 ecological reality of this rapidly growing research area was unexplored. Our objective
433 here—to survey the breadth of EcoReality in the microbiome transplant literature and
434 identify key areas lacking EcoReality—was not unlike a fact-finding mission expanding
435 the map of our understanding of reciprocal host-microbiome interactions. We
436 recommend a full, extended journey into the wilds to round out the literature’s coverage
437 of the landscape of possible EcoReality. Charting all territories, from highly controlled
438 lab studies to free-ranging organisms, is necessary to fully comprehend the interplay
439 between microbiomes and their hosts.

440

441 **Acknowledgements**

442 We thank the authors, animals, and microbes of the many studies that provided the
443 peaks and valleys that we traversed in our exploration. We thank the 3 anonymous
444 reviewers for being our north star, guiding us forward. CJGG thanks his PhD supervisor,
445 Kevin McCann, for his support. Thanks go to KAP Design for designing Figures 1 & 4.
446 We thank the attendees of the Evelyn Pielou Discussion Group for inspiring this voyage.

447

448 **Funding**

449 CJGG was supported by a Natural Sciences and Engineering Research Council of
450 Canada (NSERC) CGS-D. KC acknowledges the support of NSERC, RGPIN-2018-
451 04399. MRS was supported by a NSERC Vanier CGS.

452

453 **Author contributions**

454 All authors conceived of and produced the directed review. CJGG wrote the first draft
455 and all authors contributed to editing the manuscript.

456

457 **Data accessibility**

458 The supporting information is included with this paper. The supporting information plus
459 the data, and R script for this manuscript can be found on Zenodo/Github (41). The full
460 list of transplant studies used in this article can be found in the dataset on
461 Zenodo/GitHub (41), and (10,26,31,35,42–92) in the bibliography below.

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Supporting Information

Into the wild: microbiome transplant studies need broader ecological reality

Authors

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Search terms used for methodological literature search

- Google scholar
 - (transplant* AND microbio) (gut OR fecal OR feces OR gastrointestinal OR gastro-intestin* OR faecal OR faeces OR forces OR faecal) -human -patient -"homo sapiens" -man -woman -child*
- Web of Science
 - ((TOPIC:(transplant) AND TOPIC: (microbio))AND (TOPIC: (((((((gut OR fecal) OR feces) OR gastrointestinal) OR gastro-intestin) OR faecal) OR faeces) OR foeces) OR faecal) NOT TOPIC:((((human OR patient) OR homo sapiens) OR man) OR woman) OR child*))

Table S1 Ordinal data scale (EcoReality score) for each experimental condition

Experimental Condition	Ordinal Data Scale
Taxon Match	1 = Mismatch (different species) 2 = Match (same species)
Donor Environment	1 = Lab animal host in sterile lab 2 = Lab animal host in non-sterile lab 3 = Captive bred wildlife (multiple generations bred in captivity) 4 = Wildlife brought into captivity (no generations bred in captivity) 5 = Free-ranging wildlife (capture and release)
Donor Physiology	1 = Gene knockout or disease harbouring (non-microbial) 2 = Wildtype non-diseased
Transplanted Microbiome	1 = Single strain 2 = Consortium, mixture of select strains

	3 = Whole community (no sorting or altering of community sampled for transplantation)
Transplant Method	1 = Active (microbiome sample forcefully added to recipient gut e.g., by a suppository or oral gavage) 2 = Passive (microbiome sample passively given to recipient e.g., mixed into food)
Recipient Microbiome	1 = Germ-free 2 = Antibiotic perturbed/pathologic 3 = Whole community (no experimental alteration of community)
Recipient Environment	1 = Lab animal host in sterile lab 2 = Lab animal host in non-sterile lab 3 = Captive bred wildlife (multiple generations bred in captivity) 4 = Wildlife brought into captivity (no generations bred in captivity) 5 = Free-ranging wildlife (capture and release)
Recipient Physiology	1 = Gene knockout or disease harbouring (non-microbial) 2 = Wildtype or non-diseased
Housing Conditions	1 = Housed singly (after microbiome transplantation) 2 = Co-housed (after microbiome transplantation with either other replicates in the experiment or with individuals of the same species that were not replicates. Co-housing could also have been used as the method of transplantation)

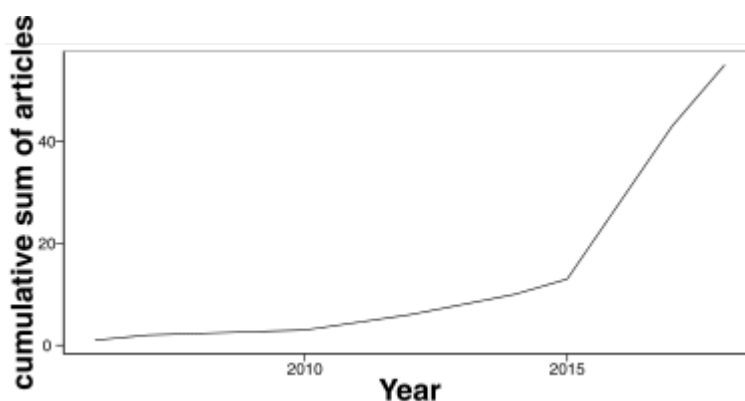


Figure S1 Cumulative sum of articles from our directed review between 2006 and 2018.

Data accessibility

The data, the above supporting information, and the R script for this manuscript are in a repository on GitHub. This repository can be cloned or downloaded straight from Github (https://github.com/cgreysongaito/Intothewild_Microbiome) or from Zenodo (<https://doi.org/10.5281/zenodo.2652255>).

Folder and file structure of Github repository (Intothewild_Microbiome)

- data
 - EcoRealTable_2019-10-09_Data.csv
- figs
 - 2019-10-09 CountAnimals.pdf – Figure 2 in manuscript
 - 2019-10-09 Eco-realityComparisons.pdf – Figure 3 in manuscript
 - 2019-10-09 Eco-realityAverageStandardOverTime.pdf – Figure 4 in manuscript
 - 2019-10-09 CumulativeSumArticles.pdf – Supporting Information Figure 1
- .gitignore – File containing files or folders that git should ignore
- IntotheWild_Microbiome_Greyson-Gaito_etal_2019.R – R script for analysis and figure creation
- SupportingInformation_Intothewild_GreysonGaitoetal.pdf – Supporting information (search terms, ordinal data scales, figure)
- LICENSE – Mozilla Public License 2.0
- README.md – Important information
- meta_transplant_microbiome.Rproj – R Project to increase ease of use

Instructions for use

- Download the whole repository (either by forking and cloning or by downloading a ZIP folder)
- In RStudio, open the project called meta_transplant_microbiome.Rproj and open the file Intothewild_Microbiome_Greyson-Gaito_etal_2019.R
 - If not using RStudio, open the file Intothewild_Microbiome_Greyson-Gaito_etal_2019.R and edit the path to the data file called EcoRealTable_2019-10-09_Data.csv to whatever path is required on your computer.
- Run the script in RStudio or however you normally run R scripts