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2	and downed woody debris
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24	Abstract. Seedling establishment is crucial for the development of self-regenerating tree
25	populations. Determinants of tree establishment vary widely and may compound in their effects.
26	Using a factorial experiment, we manipulated invasive shrubs, downed woody debris (DWD),
27	and rodent access to evaluate factors limiting the establishment of six woody species (five native
28	trees, one invasive shrub). Our results suggest these factors independently contribute to tree
29	seedling establishment. Exclusion of rodents increased establishment three-fold. Invasive shrub
30	removal (Elaeagnus umbellate; Lonicera maackii) and the presence of DWD promoted
31	establishment of two native trees (Pinus strobus; Sassafras albidum). Notably, the presence of
32	DWD halved L. maackii establishment. In identifying rodents as drivers of seed limitation, our
33	results support findings that seed additions will likely promote woody seedling establishment
34	when rodents are not abundant (e.g., low populations) or when seeds are physically or
35	chemically protected (e.g., via taste deterrents). Management plans vary in DWD retention;
36	results from our experimental cohort indicate retaining or introducing DWD promotes native tree
37	recruitment and limits invasive shrub establishment. Future studies exploring the species-
38	specific effects of invasive shrub removal and DWD amendments across multiple cohorts will
39	help determine which woody species benefit most from these management practices.
40	
41	Keywords: Tree regeneration; Seedling Establishment; Invasive Shrubs; Woody Debris; Seed
42	Predators
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47 Introduction

Sustainable forestry requires a mechanistic understanding of how the environment 48 influences tree population regeneration (Vickers et al. 2019ab, Piana et al. 2021), generating a 49 constantly evolving challenge for management as regional climates are redefined (Millar and 50 Stephenson 2015). The establishment of tree seedlings – defined as the emergence and 51 52 persistence of tree seedlings for >1 year (Clark et al. 1999) – is essential for the development of robust and self-regenerating tree populations that shape future forest structure (Hurtt and Pacala 53 1995; Norghauer and Newberry 2011; McConkey et al. 2012; Forsyth et al. 2015). Given the 54 55 fundamental role of seedling establishment in determining the dynamics of future forests (Vickers et al. 2019a; Piana et al. 2021) and the low potential or regeneration success in many 56 northern temperate U.S. forests (Vickers at al. 2019b), it is critical to identifying what and how 57 environmental factors limit or promote early establishment in these forests. 58 A primary challenge to maximizing tree recruitment is that multiple, diverse, and 59 60 interacting factors shape tree seedling establishment and survival (Goldberg 1985; Royo and Carson 2006; Piana et al. 2021; De Lombaerde et al. 2020). Competition with overstory trees 61 and vegetation in the sub-canopy can lead to light limitation, and this competition is a bottleneck 62 63 in the development of tree seeds into seedlings (Bolton and D'Amato 2011; Urgenson et al. 2012). Consumption of seeds and seedlings by animals limits juvenile tree seedling 64 65 establishment and survival (Goldberg 1985; Gill 1992ab, Crawley and Long 1995; Zwolak et al. 66 2010; Boone et al. 2019; Piana 2019), with the potential to influence the trajectory of forest community structure (Hulme and Kollmann 2005; Norghauer and Newbery 2011). Downed 67

- 68 woody debris (DWD) modifies forest floor microclimate with context-dependent and species-
- 69 specific effects on germination and seedling persistence (Harmon et al. 1986; Gray and Spies

1997; Ettinger et al. 2017; De Lombaerde et al. 2020). Complicating matters, these factors rarely 70 operate alone: competition, granivory, herbivory, and microsite limitation co-occur. For 71 example, while invasive shrubs compete for resources with tree seedlings in the forest 72 understory, the presence of these invasive shrubs also correlates with more granivore activity 73 (Dutra et al. 2011, Guiden and Orrock 2019), prolongs granivore foraging time (Mattos and 74 75 Orrock 2010), results in greater tree seed removal (Bartowitz and Orrock 2016), and alters tree seed consumption and caching near DWD (Guiden and Orrock 2017). While consensus is 76 developing regarding how these ecological factors may *independently* alter tree seed survival and 77 78 seedling establishment (e.g., Bartowitz and Orrock 2016; Ettinger et al. 2017), our understanding of regeneration in northern temperate forests will be furthered by empirical evidence examining 79 how these ecological factors may synergistically – or antagonistically – interact to shape juvenile 80 tree survival. 81

Management is most effective when foresters can act according to the relative importance 82 of constraints on seedling establishment (Webster et al. 2018). To combat competition with 83 invasive shrubs, for example, forests in the midwestern and eastern United States often undergo 84 understory clearing and managers use follow-up control measures (e.g., herbicides) to eliminate 85 86 these shrubs, restore native forest structure, and promote target seedling establishment (Hartman and McCarthy 2004; Shields et al. 2015; Ward et al. 2018). Foresters may employ unique 87 88 sowing strategies (e.g., broadcast seed timing, sowing depth) or introduce physical and chemical 89 barriers to overcome seed limitation due to consumption by animals (Willoughby et al. 2011; Leverkus et al. 2015; Löf et al. 2019). Management strategies often recommend introducing or 90 91 retaining heterogeneity in the forest landscape (e.g., snags, DWD, canopy thinning, gap 92 generation) to increase the likelihood of suitable microclimates for tree seedling establishment

and survival (Gray and Spies 1997; Bolton and D'Amato 2011; Ettinger et al. 2017; De 93 Lombaerde et al. 2020). Employing these techniques requires knowledge regarding whether and 94 how environmental factors interact to shape juvenile tree survival and performance. A strong 95 interaction between two factors, for example, may require a context-specific management 96 strategies that addresses the joint effects. Recommendations to retain or introduce DWD on the 97 98 forest floor may increase suitable microclimate for seed germination and protection of saplings from ungulate browsing (Whyte and Lusk 2019) but adding DWD concurrently introduces 99 refuge for small mammal seed predators that reduced tree seed survival in certain forest structure 100 101 (e.g., Schnurr et al. 2004; van Ginkel et al. 2013; but see Ettinger et al. 2017). Experiments are needed that evaluate how different forest understory structures shape the independent and 102 interactive effects of granivores and DWD on tree seedling establishment in temperate mixed 103 104 deciduous forests.

We use experimental removal of invasive shrubs (i.e., a manipulation of forest understory 105 structure), manipulations of DWD, and exclusion of small mammal granivores to quantify how 106 invasive shrubs, DWD, and rodents may act, alone or in concert, to modify tree seedling 107 establishment in a mixed deciduous northern temperate forest. We focused our study on species 108 that are integral to forest development and management in upper Midwest forests, examining 109 how the removal of invasive shrubs, Lonicera maackii [Amur's Honeysuckle] and Elaeagnus 110 umbellata [Autumn Olive], affects the recruitment of Acer rubrum [Red Maple], Pinus strobus 111 112 [Eastern White Pine], Quercus rubra [Northern Red Oak], Sassafras albidum [Sassafras], and *Tsuga canadensis* [Hemlock]. *L. maackii* and *E. umbellata* are invasive shrubs that negatively 113 affect native trees (Catling et al. 1997; Orrock et al. 2015), and they may limit tree recruitment 114 115 via competition, by changing small mammal granivory, or both (Orrock et al. 2015). Given that

studies have regularly reported more granivory in midwestern U.S. forests invaded by non-native 116 shrubs (Orrock et al. 2010; Bartowitz and Orrock 2016) and rates of tree seed consumption can 117 be greater near DWD (Schnurr et al. 2004; van Ginkel et al. 2013), our experiment is explicitly 118 designed to determine how invasive shrubs in a midwestern U.S. mixed deciduous forest affect 119 how DWD and seed-eating rodents interact to influence tree seedling recruitment. By evaluating 120 121 multiple, interactive factors that affect tree recruitment, our study will help managers identify conditions in mixed-deciduous forest where recruitment from seed should be highest; our study 122 will also provide an example of whether common management tools (e.g., invasive shrub 123 removal, leaving DWD following harvest) may yield maximum benefit in conjunction with 124 broadcast seed sowing in this forest context. 125

126

127 Methods

128 Field Site

We conducted replicated manipulations of invasive shrub presence at Fish Lake 129 Environmental Education Center (FLEEC), a 240-acre mixed hardwood forest property operated 130 by Eastern Michigan University. Western portions of FLEEC property were historically in 131 132 agricultural and plantations until the property was acquired by the Eastern Michigan University in 1965, when cultivation of agricultural and wooded lands ceased. The current overstory 133 includes *Quercus rubra* and other hardwoods (*Carva ovata*, *Acer* spp.) with associated conifers 134 135 (e.g., Pinus strobus, Pinus resinosa). Invasive woody shrubs encroachment is prevalent in the post agricultural portion of the FLEEC property with the dominant introduced shrubs being E. 136 umbellata and L. maackii. 137

138

In May 2018, we delineated fourteen (14) 20×20 m plots and then stratified and paired

139	these plots along a west to east gradient to generate seven blocked plot pairs; the minimum
140	distance separating plots was 50 meters. One plot in each block was randomly assigned to have
141	the invasive shrub layer mechanically removed and chemically controlled ("Invaders
142	Removed"), the invasive shrub layer was left intact for the other plot in each block ("Invaders
143	Present"). Prior to invasive shrub removal, we generated size-class distributions of the two
144	dominant invasive shrubs, <i>L. maackii</i> and E. <i>umbellata</i> , in 5×5 m sub-plots randomly
145	positioned within the larger 20 m \times 20 m plot. For 12 of the 14 sites, we measured
146	Photosynthetically Active Radiation (PAR) transmittance (1.25 m height) using a Decagon
147	Ceptometer (Decagon Devices, Pullman, WA) at 2 m meter intervals along a linear transect
148	running diagonally through the center of each plot. Ambient PAR measurements were taken
149	simultaneously with a handheld PAR sensor (Decagon Devices, Pullman, WA) to generate
150	estimates of PAR interception at each plot (see Figures S1A and S1B).
150	estimates of 171K interception at each plot (see 1 igures 517 and 51D).
150	In June 2018, we imposed our "Invaders Removed" treatment. All invasive stems within
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161 *Native tree and exotic shrub seedling establishment*

162	To test the effects of downed woody debris on woody seedling establishment, each plot
163	was divided in half (i.e., "half-plot") and DWD treatment manipulations ("DWD absent" or
164	"DWD present") were randomly assigned to one half of each plot. To generate the DWD absent
165	treatment, DWD was removed from a 2.5 m \times 5 m section from the center of the assigned plot
166	half. The removed DWD was relocated to the center of the other plot half and deposited in a
167	loose network covering approximately 2.5 m \times 5 m with a maximum height of 30 cm. We made
168	significant effort to match experimental DWD deposit structure to nearby natural DWD
169	ammalgations, with particular emphasis on mainitaining comparable heights and stem density
170	between natural and artificial DWD structures. We standardized our DWD structure as
171	described in similar studies (van Ginkel et al. 2013; see Fig. S2 for representative photos of this
172	treatment) and focused on using DWD only found within the plot to ensure we did not modify
173	the composition (i.e., source species) or the total amount of DWD found on a whole plot.
174	Consistent with results reported in similar systems, ancillary experiments (Supplemental
175	Data 3) indicate that representative granivore activity is likely lower on shrub-cleared plots
176	relative to invaded plots (Fig. S3A; Connolly et al. 2021) and that the rate of tree seed removal is
177	slower on plots with invasive shrubs removed relative to shrub invaded plots (Fig. S3B). To test
178	the effect of seed predators on woody seedling establishment, we nested two types of exclosures
179	within the center of each DWD half-plot treatment (four total exclosures per plot). One of the
180	exclosures in each half-plot pair excluded small mammal seed predator entry ("Rodents
181	Excluded") whereas the other exclosure permitted seed predator entry through holes cut in the
182	side of the exclosure ("Rodents Permitted"). In August 2019, we embedded each exclosure 25
183	cm deep in the mineral soil to prevent small mammals burrowing into the cages. Squares of
184	hardware cloth were secured as lids. To generate the seed predator access treatment, we

randomly chose one of the two exclosures on each half-plot half and cut two 5×5 cm openings on opposite sides of the exclosure to permit small mammals entry. All exclosures were constructed of 1×1 cm hardware cloth secured in a ring with wire cage clips so that the diameter of each exclosure was 15 cm (0.017 m² surface area); the small footprint and large mesh size of our exclosure was selected to minimize the effects of exclosure-mediated microclimate modifications on tree seedling establishment (Evans et al. 2018).

On 1 November 2019, seeds of six woody species common on the FLEEC property were 191 sown into each exclosure: Sassafras (Sassafras albidum), Red Maple (Acer rubrum), Canadian 192 193 Hemlock (Tsuga canadensis), Northern Red Oak (Quercus rubra), Eastern White Pine (Pinus strobus), and Amur's Honeysuckle (Lonicera maackii). Seeds of S. albidum, A. rubrum, T. 194 canadensis, and Q. rubra were purchased from a commercial vendor (Sawyer Nursery Inc., 195 Hudsonville, Michigan, USA) focusing on seed accessions from upper Midwestern U.S. states. 196 Pinus strobus seeds were provided by the Michigan Department of Natural Resources and were 197 198 sourced from collected accessions from southern Michigan. Lonicera maackii seeds were sourced from mature fruit collected from parent plants on the FLEEC property. Prior sowing all 199 S. albidum, A. rubrum, T. canadensis, P. strobus, and L. maackii seeds were first sorted using 200 firmness, and viability of subsamples was confirmed with cut tests (Karrfalt 2008). Quercus 201 rubra seeds were sorted by the float test, and a subset of the acorns were cold stratified for 21 202 days and their viability was confirmed via a germination test. These tree species are ideal for 203 204 this study because 1) all study species are found on and around the FLEEC property (Springer and Parfitt 2010, Hanes and Bunker, unpublished data), 2) represent common overstory species 205 206 found in central and eastern Michigan (Petrides and Wehr 1998), and 3) post-dispersal seed 207 predators forage on these woody species (e.g., Mattos et al. 2013, Bartowitz and Orrock 2016,

Guiden and Orrock 2017, Chandler et al. 2020) indicating that we are likley to detect
relationships between seed removal and seedling establishment if these responses are correlated
with each other. Ten seeds of each species (only five *Q. rubra* acorns) were sown into each
exclosure to mimic low to moderate natural woody plant seed density in the soil (Leckie et. al.
2000). Plots were monitored until the cessation of the experiment on 27 May 2021, when we
counted final seedling establishment.

214

215 Data Analysis

We used generalized linear mixed models using Template Model Builder with a binomial 216 distribution to estimate how the independent and interactive effects of invader removal, 217 manipulation of DWD, and seed predator access influenced the proportion seedling 218 establishment 19 months after sowing (i.e., our final seedling establishment estimates). Little to 219 no Q. rubra, A. rubrum, and T. canadensis established across all experimental units (established 220 seedlings: *Q. rubra*, n=3; *T. canadensis*, n=2; *A. rubrum*, n=0), precluding analysis of these 221 species. We ran separate models for the species with establishment (P. strobus, S. albidum, L. 222 *maackii*) and the random error structure modelled in our analyses accounted for the split-plot 223 224 design of experiment; i.e., invasive cover manipulated within block and CWD manipulated within invader treatment plot. Establishment of S. albidum was almost exclusively in units 225 excluding granivores, which precluded full model convergence. Consequently, we reduced the 226 227 structure of the model evaluating S. albidum establishment to include invader removal, DWD manipulation, and granivore access as main effects and a single interaction term between invader 228 229 removal and DWD manipulation. We used R (R Core Team 2022) and associated packages for 230 all statistical analysis and graphics generation: "ggplot2" (Wickham 2016), "car" (Fox and

Weisberg 2019), "glmmTMB" (Brooks et al. 2017), "ggpattern" (FC and Davis 2022), and
"emmeans" (Lenth 2022).

233

234 Results

Across all treatment levels, ~7.7% of sown *L. maackii* seeds established as seedlings. 235 236 Excluding granivores from sown L. maackii seeds resulted in threefold greater L. maackii seedling establishment (Table 1; Fig. 1). DWD presence resulted in ~50% less L. maackii 237 establishment, but invasive shrub cover did not affect L. maackii establishment (Table 1). 238 239 Approximately 5.0% of sown P. strobus seeds established as seedlings. Excluding granivores from sown P. strobus seeds increased P. strobus seedling establishment fourfold (Fig. 240 1, Table 1). Across all treatment levels, the presence of DWD resulted in 2.6 times greater P. 241 strobus seedling establishment (Table 1). P. strobus seedling establishment tended to greater in 242 plots with invaders removed, but the differences in means between the invasive shrub removal 243 treatments (presence versus removal) was not statistically significant at a Type I error equal to 244 0.05 (Table 1). 245 Granviore exclusion resulted in significantly greater S. albidum seedling establishment 246 247 (Table 1). Only 0.4% of S. albidum seedlings - one seedling of 280 sown S. albidum seeds -

established in plots that permitted granivore access, but $\sim 10\%$ of total sown *S. albidum* seeds (29

seedlings total) established in plots that excluded granivores (Fig. 1). Removal of invasive

shrubs and the addition of DWD independently increased average *S. albidum* seedling

establishment (Table 1). Fourfold more *S. albidum* seedlings established in plots where invasive

shrubs were removed than plots where invasive shrubs were left intact; the presence of DWD

resulted in over threefold more *S. albidum* seedling establishment than when DWD was removed

254 (Fig. 1).

255

256 Discussion

Sustainable forest management ultimately requires identifying the factors affecting the 257 establishment of trees from seeds (Webster et al. 2018; Piana et al. 2021). By monitoring seed 258 259 and seedling fate for 19 months, we gain insight into how multiple factors (i.e., invasive shrubs, native animals, and downed woody debris) can affect the native tree and invasive woody shrub 260 establishment in a mixed deciduous forest. Our study provides insight into the hierarchy of 261 262 establishment limitations encountered by dispersed seeds of woody plants during our study period: 1) invasive shrubs in the canopy generate species-specific limitations on native tree 263 seedling establishment and have little effect on invasive shrub establishment, 2) DWD promotes 264 establishment of dispersed native seeds and limits invasive shrub establishment, and 3) seed and 265 seedling limitation by granivores results in reductions in native woody plant establishment with 266 similar, but muted, effects on invasive shrub establishment. Importantly, we did not find strong 267 interactions among these three factors during our study, suggesting that in certain forest contexts 268 managers may consider each factor independently when planning management. Our work has 269 two primary implications: 1) ecological barriers vary predictably in how strongly they affect the 270 establishment of tree seedlings in certain years, and 2) forest context prior to reforestation (e.g., 271 presence of invasive shrubs, DWD coverage, small mammal populations) could generally inform 272 273 the likelihood of successful seedling establishment from natural seed rain or from reforestation using seed additions. 274

275

276 Ecological Barriers to Tree Seedling Establishment

Invasive shrubs introduce significant interspecific competition in forest understories 277 (Gorchov and Trisel 2003; Orrock et al. 2015), but the effects of invasive shrubs on the 278 establishment of woody species may be species-specific (Fig. 1; Gorchov and Trisel 2003; 279 Urgenson et al. 2012). Several mechanisms may explain why the positive effects of invasive 280 shrub removal differed in magnitude between S. albidum and P. strobus. Native species 281 282 response to invasive shrub removal may correlate with the degree of shade tolerance: S. albidum is intolerant of shade, whereas P. strobus has intermediate shade tolerance (Burns and Honkala 283 1990). Invasive plants removal promotes rapid growth in seedlings of early seral trees, but 284 285 shade-tolerant trees can display a muted response to this same treatment (Urgenson et al. 2012). Low seed addition density may also contribute to low *P. strobus* establishment in invasive shrub 286 removal plots. Fewer potential germinants in each exclosure increases the likelihood that 287 stochastic factors may swamp signals associated with the invasive shrub treatment. Our seed 288 addition density (~ 2650 seeds m⁻²), however, aligned with the ranges reported in naturally 289 occurring temperate forest seed banks (Leckie et. al. 2000 and references therein) and limits the 290 potential to overestimate the effects seed predation. Finally, seed predator activity was 291 comparable between invasive shrub treatment plots in Fall 2019 (Fig. S3) suggesting granivory 292 may have been similar between the invasive shrub treatment plots during the year seeds were 293 sown. Our work provides a detailed examination of how one experimental cohort responds to 294 these ecological barriers, but empirical evaluations of potentially interacting factors shaping tree 295 296 seedling establishment (e.g., granivory exclusion and invasive shrub removal) may be most informative when seed additions are repeated and monitored over multiple, successive years. 297 298 Downed woody debris is a common feature of forest floors (Harmon et al. 1986). 299 Greater DWD density can correspond to greater tree seedling establishment (Schnurr et al.

2004), but the effects of DWD on seedling establishment may be species-specific. DWD halved 300 L. maackii seedling establishment and the effect of DWD on L. maackii establishment was 301 cumulatively greater when the invasive shrub understory was intact (Fig. 1). Light attenuation 302 by both canopy and understory vegetation (Fig. S1B) and DWD (Gray and Spies 1997) may limit 303 photosynthesis, pushing *L. maackii* below its physiological limits in forest interiors (e.g., LCP; 304 305 Lieurance and Landsbergen 2016), increasing soil moisture (Gray and Spies 1997, Roberts et al. 2005), and lowering seed survival by facilitating soil pathogens that grow in darker, damper soils 306 (Taher and Cooke 1975; Augspurger 1990; Orrock et al. 2012). For native woody species, 307 308 however, DWD promoted seedling establishment (Fig. 1). Native tree seed germination and seedling survival can be greater on or around DWD collections (Gray and Spies 1997; 309 O'Hanlon-Manners and Kotanen 2004; Kupferschmid and Bugmann 2005). DWD can 310 diminished air and soil temperature extremes and increase soil moisture (Roberts et al. 2005; 311 Goldin and Hutchinson 2013; Dhar et al. 2022), which can foster tree seedling survival (Harmon 312 and Franklin 1989; Gray and Spies 1997) and buffer tree seedling growth in dry years (Roberts et 313 al. 2005). Currently, it is unclear what mechanisms resulted in the divergent responses we 314 observed for native versus invasive woody species establishment, but the joint manipulation of 315 316 invasive plant cover and DWD can promote woody seedling survival in certain forest contexts (e.g., urban forests, Ettinger et al. 2017). We did not, however, detect strong interactive effects 317 between these factors in this mixed deciduous forest during our study period suggesting that 318 319 interannual climate trends or other factors (e.g., forest position along a rural-to-urban gradient; Ettinger et. al. 2017) may alter the degree to which different environmental factors shape tree 320 321 regeneration from seeds.

322

Exclusion of seed-consuming animals, predominantly white-footed mice (Peromyscus

leucopus) and eastern chipmunks (*Tamias striatus*; see Supplemental Data 3), generated a nearly 323 ubiquitous increase in woody seedling establishment. Seed consumption is a well-documented 324 barrier to tree seedling establishment (Gill 1992a; Zwolak et al. 2010), with the potential to 325 decrease basal area production and slow tree population growth (Norghauer and Newberry 2011; 326 Forsyth et al. 2015). Sown seeds may have been dispersed to caches (Vander Wall et al. 2005), 327 328 but we observed few to no seedlings emerging outside our cages on plots and none of the seedlings we did observe were clumped together suggesting germination of a cache (B. 329 Connolly, personal observation). Notably, seed consumption contributed to minimizing 330 331 honeysuckle establishment suggesting granivores can play a role minimizing invasive shrub recruitment (i.e., biotic resistance), although magnitude of granivory's effect on invasive plant 332 establishment will depend on propagule pressure (Davis 2009) and the extent to which habitats 333 differ in invasibility (Connolly et al. 2014). Granivory undoubtedly shaped seedling 334 establishment in this mixed deciduous forest during this study period, underscoring the 335 importance of long-term monitoring of tree seed additions to determine how exclusion of 336 granivores translates to forest productivity. 337

Limitation by inadequate microsite conditions, non-target biotic agents, or adverse 338 edaphic conditions likely influenced test species establishment and precluded significant 339 establishment three of our five native tree species. For example, when rodents were excluded, P. 340 strobus and S. albidum establishment was low when invasive shrubs were present and DWD was 341 342 absent (Fig. 1), suggesting that adverse environmental conditions (e.g., greater temperature extremes, low soil water content) drove low establishment for these two species on invader 343 present plots. Importantly, the presence of DWD appears to ameliorate these poor establishment 344 345 conditions, resulting in a more than three-fold increase in establishment on shrub invaded plots

for these species. *Ouercus rubra* acorns only remained intact in exclosures that excluded seed 346 predators; however, only three of the 180 intact acorns in rodent exclusion plots emerged as 347 seedlings despite >80% viability at the experiment's sow date. This result suggests that the 348 environmental conditions manipulated in our study (e.g., invasive shrub cover, DWD 349 manipulation) did not promote O. rubra establishment, and other environmental factors not 350 351 directly manipulated in our study (e.g., edaphic conditions, invertebrate predation) limited native tree recruitment. For example, Raynal et al. (1982) demonstrate that low substrate pH can limit 352 the Acer rubrum and Tsuga canadensis germination, although soil pH estimates for our study 353 354 plots (Supplemental Data 4) did not reach the low pH levels used in their study. Winter conditions or invertebrate granivores and herbivores can also drive low seed survival for A. 355 rubrum and T. canadensis in Midwestern U.S. forests (Chandler et al. 2020; Guiden and Orrock 356 2021). For example, consumption by small invertebrates (e.g., earthworms, mollusks) that could 357 still access the added tree seeds in rodent exclosures may have compensated for experimentally 358 excluded small mammal seed predators (Harper 1977, Cassin and Kotanen 2016), by minimizing 359 post-emergence woody plant seed and seedling survival and effectively forestalling recruitment 360 for A. rubrum and T. canadensis (B. Connolly, unpublished data). Our study demonstrates that 361 362 tree seedling establishment is the product of multiple and diverse environmental factors and while our study helps rank predominant ecological filters to tree seedling establishment, other 363 364 environmental factors (e.g., climate extremes) will also contribute to limitation and may need to 365 be accounted for in management approaches for different forest contexts.

366

367 Management Strategies Targeting Tree Seedling Establishment

368 Our work provides empirical support that certain forest management practices increase

369 tree seedling establishment. Broadcast seed sowing, for example, may be an effective means of promoting tree seedling establishment, but only if steps are taken to minimize the effects of 370 herbivores and competition with other vegetation (Willoughby et al. 2004; Willoughby and 371 Kinks 2009; Overdyck et al. 2013; Löf et al. 2019). Seed addition often promotes seedling 372 establishment and managers may consider how the timing of seed additions, such as during 373 374 different seasons (Radvanyi 1970; Tilki and Alptekin 2006), will minimize the effects of granivores on tree seedling emergence. Rodent population abundance often varies dramatically 375 from year to year (Sullivan et al. 2023), such that adding seeds during times of low rodent 376 377 abundance, or when rodents are sated due to masting events, may increase seedling recruitment (Schnurr et al. 2004). Coating broadcast seeds in repellents may similarly be an effective means 378 of deterring seed consumers (Willoughby et al. 2011); naturally occurring compounds (e.g., 379 capsaicin) may be an effective means of deterring seed predators and some in restoration systems 380 suggests the effects of seed coat deterrents can promote target plant establishment (Pearson et al. 381 2019; Lanni et al. in press). When seeds are protected from rodents or sown during a time when 382 there is likely to be less granivory, targeting seed broadcasting on or around naturally occurring 383 collections of DWD may simultaneously promote tree seedling establishment and protect 384 developing seedlings and saplings from larger herbivores (van Ginkel et al. 2013; Whyte and 385 Lusk 2019). 386

Invasive shrubs can contribute to limitation in native tree regeneration and, consequently, invasive shrub control would contribute to restoring natural tree regeneration (Ward et al. 2018). Our work corroborates the practice of invasive shrub removal as it leads to overall average increases in native seedling establishment 19 months after treatment, although this effect may be most apparent in early seral woody species such as *S. albidum* (Fig. 1). Ward et al. (2018)

demonstrated long-term effects of invasive shrub removal by reporting increases in native tree
seedling density persisting nine years after invasive shrub removal; interestingly, seedlings <30
cm tall of large-seeded native trees (e.g., *Quercus* sp., *Carya* sp., *Acer* spp., *Prunus serotina*)
recruit to higher densities in invasive shrub removal plots in this study suggesting that
environmental factors known to act strongly on larger seeded species (e.g., granivory) may be
less potent when invasive shrubs are mechanically removed.

Managing to retain or increase local aggregations of DWD will also likely facilitate 398 seedling recruitment. Our work supports the retaining of downed woody debris on plots cleared 399 400 of invasive shrubs for two reasons. First, we observed greater seedling establishment for both P. strobus and S. albidum on plots with DWD present. Second, we observed that invasive L. 401 *maackii* seedling establishment was significantly lower in the presence of DWD. The dual 402 potential to increase regeneration of desirable native trees while curtailing the recruitment of 403 invasive woody plants may be an elegantly passive strategy to restore natural regeneration or 404 promote growth of target tree species during reforestation. Site-level characteristics of DWD 405 (e.g., composition, volume, density) may alter seed predators' activity and foraging (Sullivan et 406 al. 2012; Malo et al. 2013; Guiden and Orrock 2021). Consequently, examining how differing 407 408 densities and characteristics of the DWD drive these divergent patterns across different forest contexts is an important next step to securing this practice as a management objective in shrub-409 invaded forests. 410

411

412 *Conclusions and Future Directions*

413 Managing tree seedling establishment from seed can promote sustainable forest
414 production initiatives but requires a comprehensive understanding of a forest's environmental

context to be effective. We have demonstrated that recruitment of native species to tree seedling 415 stage can be promoted through limiting the effects of granivores, targeting seed sowing on or 416 around aggregations of DWD, and removing the invasive shrub layer. While we have 417 demonstrated the potency of these environmental constraints to limit seedling establishment, 418 longer term studies are needed to track how these factors link to the generation of healthy and 419 420 robust adult trees. Describing seedling establishment is an essential first step towards regeneration, but the fate and structure of future forests will also be a function of how ecological 421 factors shape longer term tree demography and growth (Norghauer and Newbery 2011; Forsyth 422 423 et al. 2015; Royo and Carson 2022). For example, while invasive shrubs had a small effect on seedling establishment of *P. strobus*, persistent habitation underneath invasive shrubs is also 424 likely to negatively affect native tree seedling and sapling survival and growth in subsequent 425 years (Fagan and Peart 2004) suggesting this management approach may play a more significant 426 role in *P. strobus* regeneration at older life stages. Tree seedling establishment is variable across 427 time (Clark et al. 1999). Given several species failed to recruit following seed additions under 428 any experimental conditions and given we only examined the establishment of one experimental 429 cohort, our study also highlights the importance of examining how other factors such as broader 430 431 forest context (e.g., rural vs. urban forests, land-use history) or limitation by invertebrate consumers and pathogens in the soil (O'Hanlon-Manners and Kotanen 2004; Cassin and Kotanen 432 2016) may contribute to tree seedling establishment across time. Ultimately, healthy forests 433 434 must regenerate and actively, informed participation in the process of promoting tree seedling establishment is a significant step to ensure that management practice meets sustainability goals 435 on forested land. 436

437

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715 Figure Captions

- Figure 1. The effect of manipulating invasive shrub cover, downed woody debris, and
- mammalian seed predator access on the establishment (i.e., emergence and survival after 19
- 718 months) of an invasive shrub species (*Lonicera maackii* [Amur Honeysuckle]; dark gray
- columns) and two native tree species (*Pinus strobus* [Eastern White pine] and *Sassafras*
- 720 *albidum* [Sassafras]; light gray columns); columns represent means \pm SE.

Table 1. Results from generalized linear models testing for the effect of rodent access, invasive shrub removal, manipulations of downed woody debris, and all possible higher order interaction terms on the proportion of *Pinus strobus* and *Lonicera maackii* seeds that become established seedlings. Rodent access was tested independently for *Sassafras albidum* establishment because nearly all establishment was in rodent-excluded treatments, precluding the testing of interactions that included this factor. Bolded values indicate the factor was statistically significant at a Type I error = 0.05.

Factor	χ^2	d.f.	<i>p</i> -value
Lonicera maackii			
Invaded (INV)	0.85	1	0.355
Downed Woody Debris (DWD)	3.88	1	0.049
Rodent Access (ROD)	11.72	1	<0.001
$INV \times DWD$	0.12	1	0.734
$INV \times ROD$	0.02	1	0.875
$DWD \times ROD$	0.05	1	0.825
$INV \times ROD \times DWD$	0.27	1	0.602
Pinus strobus			
Invaded (INV)	2.61	1	0.106
Downed Woody Debris (DWD)	3.86	1	0.049
Rodent Access (ROD)	12.04	1	<0.001
$INV \times DWD$	0.60	1	0.437

$INV \times ROD$	0.23	1	0.630
$DWD \times ROD$	0.76	1	0.384
$INV \times ROD \times DWD$	0.30	1	0.586
Sassafras albidum			
Invaded (INV)	4.94	1	0.026
Downed Woody Debris (DWD)	4.62	1	0.032
Rodent Access (ROD)	11.88	1	<0.001
$INV \times DWD$	2.47	1	0.116

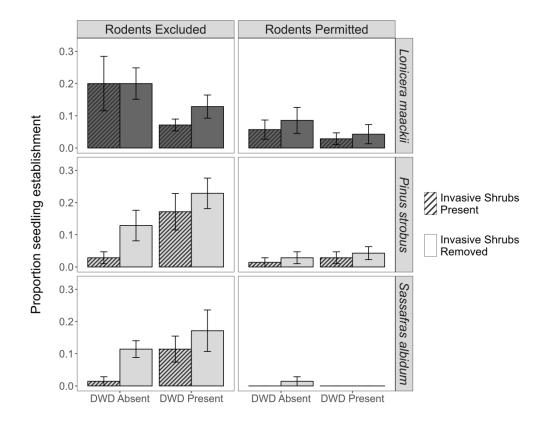


Figure 1. The effect of manipulating invasive shrub cover, downed woody debris, and mammalian seed predator access on the establishment (i.e., emergence and survival after 19 months) of an invasive shrub species (Lonicera maackii [Amur Honeysuckle]; dark gray columns) and two native tree species (Pinus strobus [Eastern White pine] and Sassafras albidum [Sassafras]; light gray columns); columns represent means ± SE.

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