

Main Manuscript for

Urban tree deaths from invasive alien forest insects in the United States, 2020-2050.

Emma J. Hudgins¹, Frank H. Koch², Mark J. Ambrose³, Brian Leung^{1,4}

¹ Dept. of Biology, McGill University

² USDA Forest Service, Southern Research Station

³ Dept. of Forestry and Environmental Resources, North Carolina State University

⁴ School of Environment, McGill University

*Corresponding author: Emma J. Hudgins.

Email: emma.hudgins@mail.mcgill.ca

Author Contributions: EJH and BL conceptualized the study. EJH performed the analyses and created the figures. EJH, BL, FHK and MJA wrote the paper. FHK and MJA provided insect and urban tree expertise and commented on realism of results. MJA manages the urban tree database.

Competing Interest Statement: The authors have no competing interests to disclose.

Classification: Biological Sciences: Ecology; Social Sciences: Environmental Sciences

Keywords: Forest ecology, Economic impact, Scenario modelling, Invasive species

This PDF file includes:

Main Text
Figures 1 to 4
Tables 1 to 1

1 Abstract

2 With 70% of the global population in urban centers, the 'greening' of cities is central to future
3 urban wellbeing and livability. Urban trees can be nature-based solutions for mental and physical
4 health, climate control, flood prevention and carbon sequestration. These ecosystem services
5 may be severely curtailed by insect pests, which pose high mortality risks to trees in urban
6 centers. Until now, the magnitudes and spatial distributions of mortality risks were unknown.
7 Here, we combine new models of street tree populations in ~30,000 United States (US)
8 communities, species-specific spread predictions for 57 invasive insect species, and estimates of
9 tree death due to insect exposure for 48 host tree genera. We estimate that an additional 1.4
10 million street trees will be killed by insects from 2020 through 2050, costing an annualized
11 average of US\$ 30M. However, these estimates hide substantial variation: 23% of urban centers
12 will experience 95% of all insect-induced mortality, and 90% of all mortality will be due to emerald
13 ash borer (*Agrilus planipennis*, EAB). We define an EAB high-impact zone spanning 902,500km²,
14 largely within the Midwest and Northeast, within which we predict the death of 98.8% of all ash
15 trees. "Mortality hotspot cities" facing costs of up to US\$ 13.0 million each include Milwaukee, WI,
16 Chicago, IL, and New York, NY. We identify Asian wood borers of maple and oak trees as posing
17 the highest future risk to US urban trees, where a new establishment could cost US\$ 4.9B over
18 the same time frame.

19 Significance Statement

20
21 US urbanization levels are already at 82% and are growing, making losses of ecosystem services
22 due to urban tree mortality a matter of concern for the majority of its population. To plan effective
23 mitigation, managers need to know which tree species in which parts of the country will be at
24 greatest risk, as well as the highest-risk insects. We provide the first country-wide, spatial
25 forecast of urban tree mortality due to invasive insect pests, including forecasts for each host tree
26 and each insect species in each US community. This framework identifies dominant pest insects
27 and spatial hotspots of high impact. Further, these findings produce a list of biotic and
28 spatiotemporal risk factors for future high-impact US urban forest insect pests.
29

30 Main Text

31 Introduction

32
33
34 Previous analyses suggest that impacts associated with urban trees are likely to comprise the
35 dominant share of economic damages caused by invasive alien forest insects (IAFIs) in the
36 United States (US) [1]. Urban tree populations include highly susceptible species like ash
37 (*Fraxinus spp.*) that are being decimated by emerald ash borer (EAB, *Agrilus planipennis*) [2]. To
38 eliminate the potential for injury or property damage due to dead trees, infested urban trees must
39 be treated or removed [3]. Moreover, the importance of urban forests is only expected to grow.
40 While urbanization is already very high in the US (82% in 2018), it has not yet reached saturation
41 (World Bank, <http://data.worldbank.org>, UN DESA, <http://population.un.org>). At the same time,
42 there has been a push for urban 'greening' (i.e., increasing urban forest canopy). Urban trees
43 perform many important ecosystem services, including lowering cooling costs [4], buffering
44 against flooding, increasing air quality, carbon sequestration, improving citizens' mental and
45 physical health outcomes, and creating important habitat [5,6]. The high tree mortality risk posed
46 by IAFIs can greatly diminish these myriad benefits.

47 While IAFI life histories differ, they are known to be transported long distances by
48 humans [7], potentially with similar drivers across entire dispersal pathways [8]. Thus, the
49 creation of a pathway-level damage estimate can provide insight into the benefit of limiting future
50 spread via these pathways (e.g. through quarantines, highway checkpoints to limit firewood

51 movement). Past estimates of IAFI damage have been important in providing support for
52 phytosanitary measures such as ISPM15 [10], a wood packing material treatment protocol,
53 whose adoption is growing worldwide [11]. A previous pathway-level estimate for the cumulative
54 cost of all US IAFIs was performed a decade ago and had substantial data limitations [1]. Since
55 then, contemporary advances allow direct estimates of spread for every IAFI species as well as
56 host prevalence and IAFI-induced mortality for every tree species in every community across the
57 United States. This allows not only the estimation of country-wide IAFI damages, but also IAFI
58 and host-specific damages and their spatial distribution. Further, we can examine the impact of
59 tree mortality dynamics on cost dynamics, and derive better risk assessments of not-yet
60 established pests, based their functional traits and host distributions.

61 In this paper, we synthesized four subcomponent models of IAFI invasions: 1) a model of
62 57 IAFI species' spread, 2) a model for the distribution of all urban street tree host genera across
63 all US communities, 3) a model of host mortality in response to IAFI-specific infestation for all
64 urban host tree species, and 4) the cost of removing and replacing dead trees, to provide the best
65 current estimate of the damage to street trees, including explicit estimates for all known IAFIs
66 across all major insect guilds (Fig. S1-S2).

67

68 Results

69

70 Urban tree pest exposure

71 Total tree abundance models were predictive with some outliers (Appendix S1, Fig. S4, small
72 trees: $R^2 = 0.78$, medium trees: $R^2 = 0.58$ large trees: $R^2 = 0.42$). Removing the outliers changed
73 the R^2 to 0.76 for small trees, 0.76 for medium trees, and 0.58 for large trees. Our genus-level
74 abundance models were strong but became slightly weaker for rare genus - size class
75 combinations (Fig. 1, overall R^2 for all genera of small trees: $R^2 = 0.93$, medium trees: $R^2 = 0.93$,
76 large trees: $R^2 = 0.92$). While relationships were variable across genera, the genera that were fit
77 most poorly did not make up a large proportion of predicted trees, and none were below $R^2 = 0.25$
78 (Fig. S5).

79 We tested four model types (global BRT, global GAM, customized BRT, or customized
80 GAM) to fit 1) genus-level tree presence/absence and 2) genus-level abundance models (Fig.
81 S1). The optimal genus-level fitting approach differed across genera depending on diameter
82 class, prevalence of genera, and whether presence/absence or tree abundance was the
83 response variable (Table S2). Generally, rarer genera were better fit by global BRT and GAM
84 models, which utilized information from all other species while common species were better fit by
85 customized models (Fig. S6). According to our models, while subject to regional variation, the
86 population of street trees is mostly made up of maple (*Acer*) and oak (*Quercus*), with substantial
87 ash (*Fraxinus*, Fig. S7).

88 We analyzed street trees separately from residential and community trees. Predicted
89 street tree exposure (measured as the number of predicted susceptible trees in Fig. 2a * IAFI
90 relative propagule pressure in Fig. 2b, [8]) across all tree types from 2020 to 2050 was generally
91 high in the eastern US, and only sporadically high across the western US (Fig. 2c). Predicted
92 street tree exposure was highest among maples (*Acer* spp., 25.6M predicted exposed trees),
93 oaks (*Quercus* spp., 5.9M), and pines (*Pinus* spp. 3.4M). The greatest number of trees were
94 predicted to become exposed to Jose scale (*Quadraspidiotus perniciosus*, 7.3M), Japanese
95 beetle (*Popillia japonica*, 6.7M), calico scale (*Eulecanium cerasorum*, 6.4M), San. Among
96 residential and community trees, exposure was greatest among maples, oaks, and *Prunus* spp.
97 (1.7B, 1.1B, 707M, respectively), and the most frequently predicted IAFI encounters were with the
98 same three species (Japanese beetle, San Jose scale, and calico scale).

99

100 Host tree mortality

101 The best-fitting mortality model indicated that most IAFIs fall in the low severity groups. Within all
102 severity groups, the majority of IAFIs were at the low end of severity (Fig. 3, full results in
103 Appendix S2). We define the term 'mortality debt' as the time period between an IAFI initiating
104 damage within a community and reaching its estimated asymptotic host mortality within that

105 community (see *Methods*). In our most likely mortality debt scenario (i.e., 10-year scenario for
106 borers, 50-year scenario for defoliators, 100-year scenario for sap feeders), we estimated a
107 mortality level of 0.7-2.5% beyond expected natural mortality of street trees by 2050, where our
108 most likely scenario fell on the higher end of this range (Table 1). Predicted street tree death
109 varied by a factor of four based on the mortality debt scenario, with longer debts leading to lower
110 total mortality between now and 2050 (Table 1). This was because in longer mortality debt
111 scenarios, trees experience mortality in the years 2020-2050 from IAFIs that initially established
112 in their communities in 2000 (50yr debt) or 1950 (100yr), but our highest impact IAFI (EAB) can
113 only begin to cause mortality after 2002 in any scenario. Sensitivity was driven largely by wood
114 boring species, as demonstrated by the sensitivity of mortality estimates to their mortality debt
115 scenarios (“Vary Borers” row, Table 1). We also found that longer mortality debts lead to a
116 smoother cost curve, or costs that do not vary much due to more consistent host mortality rates
117 (Fig. 4).

118 Spatially, future damages will be primarily borne in the Northeast and Midwest, driven by
119 EAB spread (Fig. 2d). We predict that EAB will reach asymptotic mortality in 6747 new cities,
120 which means that 98.98% of its preferred *Fraxinus* spp. hosts will die. Thus, the mortality is
121 predicted to be concentrated in a 902,500km² zone encompassing many major Midwestern and
122 Northeastern cities (Fig. S10). This mortality is also predicted to result in a 98.8% loss of all ash
123 street trees within this zone. Over 230,000 ash street trees are predicted to have died before
124 2020, and there are a further 69 cities where EAB is predicted to reach asymptotic mortality within
125 10 years of 2050 (i.e., 98.8% ash mortality by 2060). Due to the restricted range of forest ash
126 relative to urban ash, we predict that 68% of ash trees and 76% of communities containing street
127 ash will remain unexposed to EAB in 2060. Furthermore, at-risk ash trees are unequally
128 distributed. We projected the highest risk close to the leading edge of present-day EAB
129 distributions, particularly in areas predicted to have high ash densities. The top “mortality hotspot
130 cities”, where projected added mortality is in the range of 5,000-25,000 street trees, include
131 Milwaukee, WI, the Chicago Area (Chicago/Aurora/Naperville/Arlington Heights, IL), Cleveland,
132 OH, and Indianapolis, IN (Fig. 2d). Cities predicted to have high mortality outside of the Midwest
133 include New York, NY, Philadelphia, PA, and Seattle, WA – communities with high numbers of
134 street trees and high human population densities, which attract EAB propagules within our spread
135 model. The states most impacted by street tree mortality match these patterns, where the highest
136 mortality is predicted for Illinois, New York, and Wisconsin.

137

138 Cost estimates

139 We estimated annualized street tree costs across all guilds to be between US\$29-33M per year in
140 our most likely scenario (mean = \$30M, Table 1). Roughly 90% of all costs across the entire US
141 were due to EAB-induced *Fraxinus* spp. mortality. The total cost associated with street tree
142 mortality in the top ten hotspot cities was estimated at \$50M from 2020 to 2050, with \$13M in
143 Milwaukee, WI alone.

144 The ranking of feeding guild severity was relatively robust across mortality debt
145 scenarios, in spite of the potential for differences due to the interaction of IAFI-specific spread
146 and mortality debt dynamics. Costs were higher for longer mortality debt scenarios for borers,
147 peaked at intermediate debt for defoliators, and peaked at the longest debt for sap feeders.
148 These patterns were due to the relative rates of historical and contemporary range expansion of
149 more impactful IAFIs (i.e. high impact borers have more rapid recent range expansion, while
150 contemporary high impact defoliator expansion is slow compared to 50 years ago). Borers were
151 predicted to be the most damaging feeding guild (\$8M-28M mean annualized street tree
152 damages across scenarios), and EAB was consistently the top threat. Defoliators were predicted
153 to be the second most damaging feeding guild in the next 30 years (means = \$0.8M-\$1.4M), in
154 spite of more widespread hosts than wood borers, due to lower asymptotic mortality levels.
155 Defoliators had a 1-2 order of magnitude lower cost than wood-boring species, but again showed
156 consistency in which species were the top threats within the guild. Consistent with previous work
157 in [1], European gypsy moth had the highest cost of all defoliators, followed by Japanese beetle
158 and cherry bark tortrix (*Enarmonia formosana*). The sap-feeding group accrued the lowest costs

159 in the next 30 years due to their lower asymptotic mortality and rarer street tree hosts (mean =
160 \$0.2M-1.1M). Hemlock woolly adelgid (*Cryptococcus fagisuga*) was the highest impact sap
161 feeder, followed by oystershell and elongate hemlock scale insects (*Lepidosaphes ulmi*, *Fiorinia*
162 *externa*). Total costs were only notably sensitive to borer mortality debt scenario misspecification
163 (Table 1), which is promising, given our certainty of the shorter scenario for EAB.

164

165 Potential impacts to non-street trees

166 Mean added mortality (i.e. above background rates) for residential and non-residential community
167 trees in the most likely scenario was 1.0% (13.3M residential and 72.1M non-residential trees,
168 Table S10). While recognizing that non-street tree management will likely be more variable, to
169 provide a rough estimate, we assumed that non-street trees would be managed in the same way
170 as street trees (i.e. removal and replacement of dead trees). In this scenario, added mortality
171 would incur an estimated annualized cost of \$1.5B for non-residential trees and \$356M for
172 residential trees. Further, a disproportional amount of the total damages (91% of the mortality to
173 residential non-residential community trees) is expected to be felt in the aforementioned hotspot
174 zone, with 12.1 million residential and 65.9 million non-residential community trees expected to
175 be killed. Given the relatively limited data, and the difference in potential management behaviour
176 for these trees, we caution against overinterpretation of these results.

177

178 Novel IAFI risk forecast

179 Our framework allowed us to identify the factors leading to the greatest impacts for IAFIs already
180 known to have established in the United States. We were able to identify the most common urban
181 host trees, the sites facing the greatest future IAFI propagule pressure, and the IAFI-host
182 combinations with the greatest mortality. However, this approach can also be synthesized with
183 IAFI entry scenarios to understand potential impacts of novel invasive IAFIs. To illustrate the
184 utility of this framework predictively, we have provided a checklist of risk factors in Table S11 and
185 future spread simulations in Table S12 and Fig. S12. We show that entry via a southern port (e.g.
186 the Port of South Louisiana) would lead to the greatest number of exposed trees. Further, an
187 EAB-like borer of oak and maple trees could kill 6.1 million street trees and cost \$4.9B over the
188 next 30 years.

189

190

191 **Discussion**

192

193 While previous analyses have indicated that urban trees are associated with the largest share of
194 economic damages due to IAFIs [1,13,14], until recently, data did not exist on the urban
195 distribution of host trees [15], the spread of IAFIs [8], nor the mortality risk for hosts due to
196 different IAFIs [16]. With these new models, it is now possible to forecast where and when IAFIs
197 will have the most damages across the US. Our analysis suggests an overall added mortality of
198 between 2.1-2.5% of all street trees, amounting to \$US 30M per year in management costs.
199 However, the most interesting and potentially useful element was our ability to forecast hotspots
200 of future forest IAFI damages, including a 902,500km² region that we expect to experience 95.7%
201 of all mortality, in large part due to a 98.8% loss of its ash street trees due to EAB. This type of
202 forecasting has been highlighted as a crucial step in prioritizing management funds [17]. These
203 data can be used by municipal pest managers to anticipate future costs, and may help motivate
204 improved spread control programs that aim to identify the potential source counties of future
205 invasions and mitigate the worst anticipated impacts (complete forecast available at
206 <http://github.com/emmajhudson/USstreetdamage>).

207

208 Beyond present IAFI risks, our integrated model can also act as a risk assessment tool
209 for street tree mortality caused by novel IAFIs (Table S11-S12, Fig. S12). While ash trees are
210 assured to be dramatically affected by EAB over the next few decades, our models suggest oak
211 and maple to be the most common street tree genera nationwide. Further, while ash species are
212 being substituted for less susceptible tree species, maples and oaks continue to be widely
planted within our street tree inventories. Therefore, IAFIs with host species spanning these

213 genera should be of heightened concern. Secondly, the timescale and magnitude of the impacts
214 of wood borers (see also [1]) make them the highest risk to street trees. We integrated these two
215 pieces of information with information on major ports of entry within the US (American Association
216 of Port Authorities 2015, <http://aapa.com/>), as well as our general model of IAFI spread [8], to
217 forecast the extent of exposed maple and oak street trees from 2020-2050 (Fig. S12, Table S12).
218 Our analyses show that entry via a southern port would lead to the greatest number of exposed
219 trees. Further, larger trade volumes between the US and Asia compared to other regions [18]
220 suggest Asian natives will be the most likely future established IAFIs. One potential candidate
221 species fitting these criteria is citrus longhorned beetle, which is an Asian wood borer thought to
222 have many potential host species within the United States, including ash, maple and oak [19].
223 The lack of strict implementation of current wood treatment protocols such as ISPM15 [20]
224 increases the susceptibility of the US to invasion and subsequent spread of this species and
225 other potentially high-risk borers.

226 Our impact estimates vary substantially based on dynamics of host mortality following
227 initial IAFI invasion, especially because of variability in the duration and functional form of
228 mortality debt. Luckily, the guild (borers) and species (EAB) whose impact on total community
229 costs are most sensitive to correct specification of the mortality debt dynamics are the ones for
230 which we are most confident. Several publications have demonstrated near-complete decimation
231 of ash stands in the decade following EAB infestation [2,21,22]. Furthermore, since total tree
232 mortality is asymptotically equivalent across all mortality debt regimes, if other feeding guilds
233 possessed 10-year mortality debt regimes, we should have been able to detect a rapid die-off of
234 their hosts as they spread, similarly to what we found for EAB (albeit scaled by their maximum
235 mortality rates). This is not the case in the literature [22].

236 With our integrated model, we also estimated economic damages, which updates the
237 decade old Aukema et al. [1] using recent advances [13,14]. Surprisingly, the previous cost
238 estimates were not that different at the country scale. The previous cost estimate separated
239 urban trees into residential and non-residential types (grouping street trees in the latter). We
240 estimate annualized costs for non-residential trees to be somewhat lower than those in [1] (\$1.3B
241 versus \$2.0B in total “Local Government expenditures”). This lower estimate is likely because of a
242 lower rate of predicted *Fraxinus* exposure to EAB (i.e., lower predicted ash abundance in areas of
243 predicted EAB spread) in non-residential areas. Interestingly, our estimate of residential tree
244 costs is roughly one third that in [1] (\$303M vs \$1.1B in total “Household Expenditures”), again
245 likely due to a (more extreme) overestimate in the nationwide prevalence of residential ash trees
246 in the previous publication.

247 Additionally, we predict that 75% of communities containing ash trees and 68% of all
248 street ash will remain untouched by EAB by 2060 because of the lack of forest ash beyond our
249 forecasted invasion extent (i.e., affecting exposure). However, in some EAB infested
250 communities, it is important to note that our street tree distributional model may overestimate the
251 tree mortality projected, due to the role of preventative cutting prior to EAB arrival, which occurred
252 in many cities across IN, IL, MI, and WI. Preventative cutting would have led to the payment of
253 tree removal costs prior to our estimation window. This is particularly likely to have inflated the
254 2020-2050 costs to communities with large street tree budgets in regions where EAB was
255 predicted to invade in the years 2010-2020 (therefore initiating mortality 2020-2030).

256 Spatially, our results show clear patterning of high threat in the eastern and central US,
257 and lower threat in the western US. This pattern is consistent with previous findings [20], and can
258 be explained by the high impacts of EAB, European gypsy moth, and hemlock woolly adelgid,
259 whose distributions are projected to concentrate further east in the short term. However, some of
260 the highest-impact non-native pathogens have emerged in the western US, and were not
261 captured in this analysis [23,24]. Western regions could also see high future risks due to the
262 polyphagous shot hole borer (*Euwallacea whitfordiodendrus*), and its insect-disease complex with
263 fusarium fungus (*Fusarium spp.*) [25]. This complex has already established in California and has
264 maple and oak trees among its many hosts.

265 While the substantial advances that emerged recently allowed us to develop a more fully
266 integrated model, we also identified data deficiencies which require additional research. A relative

267 quantification of additional sources of uncertainty is provided in Appendix S3. This cost estimate
268 is arguably a lower bound, since it only examines the cutting of dead trees. The analysis also fails
269 to account for preventative management, to fully examine non-street tree management, and to
270 assess the impacts of IAFIs that have not yet established in the United States. Furthermore, our
271 analysis assumes a complete identification of 'high impact IAFIs'. Some presently established US
272 may not yet have been identified as 'high impact', either due to lags in their impact, and/or lags
273 the detection of this impact [26], but may achieve the same level of recognition as those in [1]
274 before 2050.

275

276 Conclusion

277 We have shown that the suite of known IAFIs have the potential to kill roughly a hundred million
278 additional urban trees in the US in the next 30 years. While these numbers themselves are
279 striking, reporting only a country-level impact estimate without IAFI species, tree, and community-
280 level resolution does little to inform management prioritizations. Here, we were able to identify
281 specific urban centers, IAFI species, and host tree genera associated with the vast majority of
282 these impacts. We predict that 90% of all street tree mortality within the next 30 years will be
283 EAB-induced ash mortality, and that ~95% of all street tree mortality will be concentrated in less
284 than 25% of all communities. These estimates illustrate the gravity of IAFI infestations for
285 communities in the path of high impact invaders that are rich in susceptible hosts. Further, we
286 were able to use this framework to identify a checklist of biotic and spatiotemporal risk factors for
287 future high-impact street tree IAFIs.

288

289 **Materials and Methods**

290

291 We synthesized four subcomponent models of IAFI invasions: 1) a model of 57 IAFI species'
292 spread, 2) a model for the distribution of all urban street tree host genera across all US
293 communities, 3) a model of host mortality in response to IAFI-specific infestation for all urban host
294 tree species, and 4) a simple model of the human management response to dead host trees, to
295 provide the best current estimate of the damage to street trees (see conceptual diagram, Fig. S1).

296

297 IAFI dispersal forecasts

298 We modelled spread using the Semi-Generalized Dispersal Kernel (SDK, [8]). This is a spatially
299 explicit, negative exponential dispersal kernel model that can account for additional spatial
300 predictors in source and recipient sites. The SDK builds from the Generalized Dispersal Kernel
301 (GDK, [8]) as a starting point, using human population density, forested land area and tree
302 density in source and destination sites as moderators of spread. The SDK combines up to three
303 species-specific corrections for each species to maximize predictive ability: 1) a species-specific
304 intercept term, 2) information on an IAFI's likely initial invasion location, and 3) niche-related
305 limitations when evidenced in the literature. The SDK was applied to all 57 IAFIs believed to
306 cause some damage from [1], and projected from 2020 to 2050 (Fig. S2).

307

308 Street tree models

309 Our fitting set consisted of 653 street tree databases for US communities where street tree
310 inventory data had been collected (Fig. S3, [14]). In two communities (Tinley Park and, IL and
311 Fort Wayne, IN), preventative cutting for EAB was conducted prior to the most recent inventory
312 and was therefore accounted for within our dataset. We modelled the abundance and diameter at
313 breast height (DBH) for trees within each genus, as treatment costs are dependent on number
314 and diameter of trees [1]. We split trees into three diameter classes (small = 0-30cm, medium =
315 31-60cm, large >60cm). We first fit models for the total tree abundance of all species by diameter
316 class, and then used these total tree models to help predict genus-specific tree abundance within
317 each diameter class. Street tree inventory data are not always reliably reported to the species
318 level across municipalities, and some species are so rare in street tree inventories that it would
319 have been very difficult to develop robust species-level models, so we limited our examination to
320 the genus level. Since IAFIs may not be equally impactful to all host tree species in a genus, we

321 had to estimate the genus-level severity of each IAFI species for each IAFI-host combination. We
322 did so by estimating the species-level breakdown of each genus based on their average relative
323 proportions across our 653 inventoried communities, and assuming this distribution was
324 representative in other projected communities.

325 We modelled the total abundance of street trees in a community using boosted
326 regression trees (BRT, *gbm.step* within R package *dismo*, [27]) relating the logarithmically-scaled
327 total tree abundance within a diameter class to community-specific predictors, employing
328 environmental variables from WORLDCLIM [28] and community characteristics used in [13], and
329 sourced largely from the National Land Cover Database (NLCD,[29]), the US Census and the
330 American Community Survey (<https://www.census.gov/data.html>, Table S1). We hypothesized
331 that the age and wealth of a community would influence the types and sizes of trees planted
332 there. In our model, median home value and mean year of construction (at the block-group level)
333 as well as median household income (at the county level) were used as proxies of the age of the
334 urban tree community and the community budget for street trees. We also tested the use of
335 Poisson GAM models, but high levels of concurrency (the GAM equivalent of multicollinearity, [30])
336 amongst predictors and lower predictive performance indicated Poisson GAMs were an inferior
337 modelling structure for estimating total abundance.

338 Next, we estimated the abundance of street trees within each genus, using the same
339 climatic and demographic factors as the total tree abundance model as well as the total tree
340 abundance model output as predictors (Fig. S1). We considered two approaches: 1) Zero-inflated
341 Poisson GAMs, or 2) a two-step BRT approach. For BRT, we modeled tree presence/absence,
342 followed by tree abundance given presence (using logarithmically-scaled tree abundance and
343 back-transforming when predicting), and then combined the two models. The number of trees of
344 genus i in size class j at a particular site k was:

$$345 \quad trees_{i,j,k} = c_{i,j,k} * pred_{exist,i,j,k} * pred_{number,i,j,k} \quad (1)$$

$$346 \quad c_{i,j,k} = \frac{1}{\sum_k(pred_{exist,i,j,k} * pred_{number,i,j,k}) / \sum_k obs_{number,i,j,k}} \quad (2)$$

347 This process is similar to a zero-inflated Poisson (ziP) model [31] but does not link the
348 parameters of the binary and continuous components of the model, instead fitting them
349 separately. Because our BRT approach was built from two independent parts, we needed to add
350 a rescaling step so that the output summed to the observed counts (eqn. 2), as occurs for ziP
351 models by default [31]. We removed all highly correlated variables ($r > 0.8$) prior to fitting, and
352 refit GAMs until maximum estimated worst-case concurrency using three-knot smoother functions
353 was below 0.8 (*concurrency* function within *mgcv*,[32]).

354 We compared BRT and GAM models that were fit to all genera simultaneously (general
355 BRT/GAM models using genus-specific intercept terms) with models that were fit to each genus
356 separately (customized BRT/GAM models) (Fig. S1). Predictive power could be higher when
357 modelling all genera together if the genera respond similarly to predictors, while power could be
358 higher for individually fitted genera where environmental and community characteristic
359 relationships are idiosyncratic and where the sample is sufficiently large.

360 We chose the model that produced the strongest relationship for each genus using R^2
361 values that were relative to the 1:1 line (i.e. a normalized mean squared error, R^2_{MSE}). R^2_{MSE} more
362 correctly measures deviations between observations (y) and predictions (\hat{y}) than conventional R^2 .

$$363 \quad R^2_{MSE} = 1 - \frac{\sum(y - \hat{y})^2}{\sum(y - \bar{y})^2} \quad (3)$$

364 We removed New York, NY from the fitting set as it was likely to be a high leverage observation
365 and could have significantly changed the resulting models due to it possessing a markedly
366 different street tree genus composition from all other communities. Both the GAM and BRT
367 models were fitted using their built-in cross-validation algorithms for parameter estimation, and
368 can therefore tolerate occasional outliers with minimal effect on their parameter estimates (though
369 we have less evidence that other outliers would have changed model parameters for cities other
370 than New York, NY). Given the higher data requirements of GAMs (i.e. all parameters must be fit
371 simultaneously, rather than BRT, which can fit subsets of predictors to each tree, [33]), genus-
372 specific GAMs were not considered when data were insufficient (i.e., when only a few cities

373 contained that genus). For each genus, we used the best-fitting model to predict urban tree
374 distributions throughout the contiguous US. We used the observed number of trees rather than
375 model predictions in cities where these data were available. Alaska and Hawaii were removed to
376 match the spatial extent of IAFI spread predictions, and because urban tree genus composition is
377 likely quite different in these areas compared to the contiguous US.

378 We synthesized the previous two modelling steps, intersecting IAFI spread forecasts with
379 predicted tree distributions (using observed tree data where available), to create forecasts of tree
380 exposure, which we define as the sum of predicted density of each IAFI species, multiplied by
381 their predicted host tree abundance in each community.

382

383 Host mortality model

384 We examined the impacts of the three major feeding guilds of IAFIs [34]: Foliage feeders included
385 insects that feed on leaf or needle tissue. Sap feeders included all species that consume sap,
386 including scale insects and gall-forming species. Borers included species that feed on phloem,
387 cambium, or xylem. Across insect guilds, the logic from [1] appeared to hold: most species were
388 innocuous, but a small number caused high mortality (Table S7). In contrast, while several
389 invasive pathogens were mentioned in [14], pathogens are only reliably reported when they
390 produce noticeable (i.e. intermediate) impacts [1]. To avoid mischaracterizing their impacts, we
391 removed pathogens from the remainder of our analysis.

392 We ranked the severity of a given IAFI infestation on a particular host using a scale
393 based on observed long-term percent mortality (defined in [14], Table S7). We added two
394 additional categories to this scale to represent IAFI species missing from their database that are
395 still considered pests on a particular host in [1]. The lowest-impact IAFI-host combinations were
396 those featuring IAFIs reported as 'low impact' in [1]. These accounted for most known
397 combinations. The second lowest category featured 'intermediate impact' IAFI species from [1]
398 that did not appear as threats to a given known host in [14]. We assumed that, were these
399 species quantified by [14], their associated severities would be lower than the lowest category
400 within the authors' ranking scheme. All other IAFI-host combinations were assigned to the same
401 categories as in [14]. IAFI frequency within severity categories was normalized across the sum of
402 their known hosts so that each IAFI had equal impact on the frequency distribution (i.e.,
403 frequency summed to 1 for each IAFI). For instance, if an IAFI had 3 hosts, and had severities of
404 3, 5, and 9 on each host, we would give them a frequency of 1/3 under each bin. We fit a Beta
405 distribution to the frequency distribution of IAFIs in each of these categories using Stan [35], a
406 program and language for efficient Bayesian estimation. We chose to fit a Beta distribution
407 because proportional mortality ranged between 0 and 1. Additionally, we fit the upper limit of the
408 two lowest mortality categories and the lower limit of the highest category, as these categories did
409 not have quantified bounds, but could be ranked relative to others. We used the posterior mean
410 as the expected mortality for an IAFI in each severity category, rather than the simple midpoint of
411 the range of each category.

412 We define the term 'mortality debt' as the time period between an IAFI initiating damage
413 within a community and reaching its estimated asymptotic host mortality within that community.
414 While we had estimates of the asymptotic proportional mortality of host trees [14], we had no
415 information on the rate by which trees reach this plateau. Previous estimates have ranged from 5
416 to 100 years [1,36], so we analyzed three scenarios within this range (10, 50, 100 years). To
417 account for what is currently known about the mortality dynamics of IAFIs within each of the
418 feeding guilds, we also examined scenarios based on our most likely scenario of the duration of
419 mortality debt across IAFI feeding guilds. EAB is estimated to kill the majority of its susceptible
420 hosts in the first decade following infestation [19], while maximum mortality is estimated to take
421 closer to 100 years for hemlock woolly adelgid [1], so we used the 10 and 100-year scenarios for
422 borers and sap-feeders, respectively. A recent publication examining mortality rates in forested
423 areas suggested that European gypsy moth has a mortality rate intermediate between borers and
424 sap-feeders, so we set defoliators at 50-years [20]. Once an IAFI was predicted to infest an area,
425 we imposed a 10-year initial lag phase between IAFI arrival at a site and the initial onset of
426 damage [37,38] and then began increasing the host mortality following our mortality debt scenario

427 to the asymptotic level (defined by the host mortality model). For simplicity, we assumed
428 mortality increased by a constant fraction over time until reaching its maximum and levelling off.
429 For example, in the 50-year mortality debt scenario, if an IAFI's maximum host mortality was
430 defined as 90%, mortality would increase by 9% at each 5-year timestep for 10 timesteps until
431 90% mortality had been reached.

432 The joint impact of maximum mortality and mortality debt is best illustrated by a series of
433 examples. Estimates of street tree natural mortality range around 2.4-2.6% per year [12]. Within a
434 30-year window, this would amount to roughly 53% natural street tree mortality. Our model
435 assumes that if IAFI enters site at the beginning of this window (2020), it first undergoes a 10-
436 year time lag, and can then cause mortality in the final 20 years. The maximum level of mortality
437 induced by a borer (EAB on several *Fraxinus* spp., Category H = 98.98%), would result in 98.98%
438 additional mortality (mortality of remaining the trees that survived natural mortality) at the end of a
439 30-year window. This level of mortality would be clearly detectable above natural street tree
440 mortality. Hemlock woolly adelgid has a similar maximum mortality to EAB (Category H on *Tsuga*
441 spp.), but we have assumed that sap-feeder mortality takes 100 years to reach asymptotic levels.
442 As such, by the end of a 30-year window, only $(98.98\%/100) \times 20$ years = 19.80% of additional
443 host trees would be killed. While defoliators have shorter mortality debts, they tend to cause lower
444 mortality, making their impacts the least detectable above background mortality. For defoliators,
445 the IAFI with the greatest damage on any host is the larch casebearer, (*Coleophora laricella* on
446 *Larix laricina*, category E = 16.46%). Given a 50-year mortality debt for defoliators, the maximum
447 mortality above background rates by 2050 is $(16.46\% / 50) \times 20$ years = 6.58%. While these
448 estimates are much lower, many host trees of sap feeders and defoliators are very common, and
449 this mortality could very well be inflating the perceived background mortality rates of these host
450 trees measured in [12].

451

452 Management costs

453 As a final layer that allowed us to move from mortality estimates to cost estimates, we estimated
454 the cost of removing and replacing dead trees. We used this cost because we believe it to be the
455 minimum management response required, and because the extent and variability of preventive
456 behaviour would be much harder to estimate. However, we note that this cost does not account
457 for additional preventive cutting or any non-cutting management such as spraying or soil
458 drenching with pesticides. We assumed that cutting was a one-time 100% effective treatment
459 against IAFIs, or in other words, that newly planted trees were of different species and thus not
460 susceptible to the same IAFI species that killed the previous trees. We assumed a 2% discount
461 rate for future damages [1] and also that infestations were independent, or in other words that
462 invasion by one IAFI did not interfere with invasion by another. This is likely a fair assumption, as
463 there is minimal host sharing across IAFIs, and IAFI species each infest only a small proportion of
464 hosts at a given time interval, so there is minimal potential for species interactions [30]. We
465 assumed the same per-tree cost estimates for cutting and replacing dead trees as in [1], where
466 the cost of cutting increases nonlinearly with size class. If we assume that street trees are always
467 under the jurisdiction of local governments, the cost of removal and replacement of each tree is
468 US\$450 for small trees, US\$600 for medium trees, and US\$1200 for large trees (these costs
469 jump to an estimated US\$600, US\$800, and US\$1500 for homeowners). We reported all costs
470 incurred from 2020 to 2050 in 2019 US dollars based on a 2% discount rate relative to these
471 baseline costs. Since these baseline per-tree management costs came from a 2011 publication,
472 we converted them to 2019 dollars via the consumer price index, which amounted to an inflation
473 of 13.65% (World Bank, <https://data.worldbank.org>), though we note that the present-day costs of
474 per-tree removal may have declined with advances in technology.

475

476 Model synthesis

477 Once all subcomponent models had been parameterized, we synthesized the street tree
478 estimates, IAFI spread estimates, host mortality estimates, and removal costs to produce overall
479 cost estimates (Fig. S1). We summed the damages from 2020 to 2050 to obtain a total

480 discounted cost for this 30-year window. We then obtained annualized costs by calculating an
481 annuity over the 30-year time horizon using the following equation:

$$482 \quad \text{Annualized damage} = D \frac{\sum_{\text{time}=\text{min}}^{\text{max}} \text{Cost}_{\text{time}}}{(1-(1+D)^{\text{min}-\text{max}})} \quad (4)$$

483 Where D is the discount rate (2%). Using these forecasts, we extended the concept of
484 cost-curves from [1], which were based on frequencies of occurrences of low and intermediate
485 damaging IAFI, and explicit economic estimates of three 'poster pests'. To parameterize the cost-
486 curves in this manuscript, rather than just 3 poster pests, we estimated street tree costs for all 57
487 intermediate-impact IAFIs across the 3 major insect feeding guilds, in addition to frequencies of
488 low-impact species (Table S4.1). The summed area under each guild-specific curve can be
489 interpreted as the estimate of the total annualized cost of all IAFIs in the US to street trees. Since
490 our curves were missing only low-impact species, the total cost estimated with these approaches
491 is not appreciably different from a simple sum of the costs of the non-missing (57 intermediate)
492 species reported in text, but we included these analyses to allow for the prediction of the costs of
493 novel invaders from each guild (Appendix S4).

494 We assessed parameter uncertainty in proportional host mortality by sampling from our
495 posterior beta mortality distribution. We also used sensitivity analysis to explore the effect of
496 different mortality debt scenarios, including 1) our most likely scenario, 2) setting all guilds to 10,
497 50, or 100-year debts, and 3) varying each guild separately while holding the other two guilds at
498 their most likely scenario. While our host distribution models were based on standard modelling
499 approaches (e.g. GAM), our Bayesian formulations underlying the mortality estimates were novel
500 and needed to be tested theoretically, to ensure that parameters were identifiable, and
501 reproduced the correct behavior. See Appendix S4 for details of our theoretic analyses.

502 Potential impacts to non-street trees

503 To provide a rough estimate of non-street tree impacts, we built a model for whole-community
504 trees (i.e., street + non-street trees) from the dataset of 56 communities where genus-level
505 estimates were reported, subtracted predicted street trees from this whole community estimate,
506 and apportioned the remaining trees into residential and non-residential trees based on their
507 average fractions across all sites where land type breakdowns were provided (32 municipalities).
508 Given the relatively limited data, we caution against overinterpretation of these results.

512 **Acknowledgments**

513
514 EJM would like to thank her PhD supervisory committee members T. Jonathan Davies and
515 Patrick M. A. James for their invaluable comments, as well as the thoughtful comments and
516 questions from thesis external examiner Dominique Gravel and colleague Andrew Liebhold. EJM
517 also acknowledges the continual support and feedback from lab members Dat Nguyen, Abbie
518 Gail Jones, Charlotte Steeves, Shriram Varadarajan, and Lidia Della Venezia. This work was
519 supported by a NSERC CGS-D awarded to EJM.

521 **References**

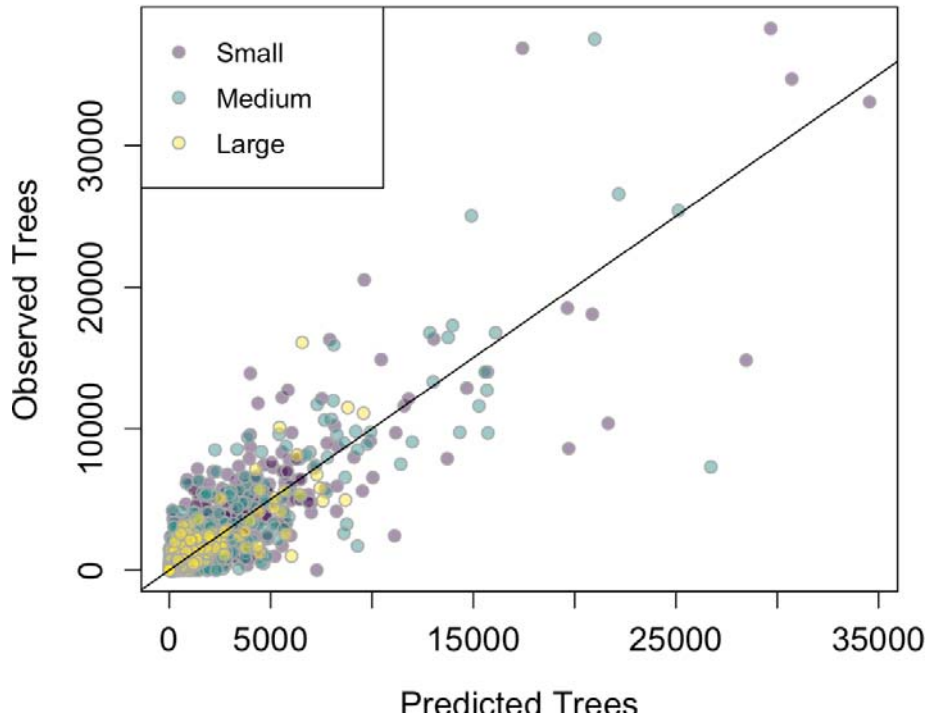
- 522 1. Aukema, J. E., Leung, B., Kovacs, K., Chivers, C., Britton, K. O., Englin, J., ... *et al.*
523 (2011). Economic impacts of non-native forest insects in the continental United States.
524 *PLoS One*, 6(9).
- 525 2. Kovacs, K. F., Haight, R. G., McCullough, D. G., Mercader, R. J., Siegert, N. W., &
526 Liebhold, A. M. (2010). Cost of potential emerald ash borer damage in US communities,
527 2009–2019. *Ecological Economics*, 69(3), 569-578.

- 528 3. Fahrner, S. J., Abrahamson, M., Venette, R. C., & Aukema, B. H. (2017). Strategic
529 removal of host trees in isolated, satellite infestations of emerald ash borer can reduce
530 population growth. *Urban Forestry & Urban Greening*, 24, 184-194.
- 531 4. Norton, B. A., Coutts, A. M., Livesley, S. J., Harris, R. J., Hunter, A. M., & Williams, N. S.
532 (2015). Planning for cooler cities: A framework to prioritise green infrastructure to mitigate
533 high temperatures in urban landscapes. *Landscape and Urban Planning*, 134, 127-138.
- 534 5. Van den Berg, A. E., Maas, J., Verheij, R. A., & Groenewegen, P. P. (2010). Green space
535 as a buffer between stressful life events and health. *Social science & medicine*, 70(8),
536 1203-1210.
- 537 6. Roy, S., Byrne, J., & Pickering, C. (2012). A systematic quantitative review of urban tree
538 benefits, costs, and assessment methods across cities in different climatic zones. *Urban
539 Forestry & Urban Greening*, 11(4), 351-363.
- 540 7. Hulme, P. E. (2009). Trade, transport and trouble: managing invasive species pathways
541 in an era of globalization. *Journal of Applied Ecology*, 46(1), 10-18.
- 542 8. Hudgins, E. J., Liebhold, A. M., & Leung, B. (2017). Predicting the spread of all invasive
543 forest pests in the United States. *Ecology letters*, 20(4), 426-435.
- 544 9. Hudgins, E. J., Liebhold, A. M., & Leung, B. (2020). Comparing generalized and
545 customized spread models for nonnative forest pests. *Ecological Applications*, 30(1),
546 e01988.
- 547 10. IPPC (International Plant Protection Convention). 2002. International standards for
548 phytosanitary measures. Rome, Italy: FAO.
549 www.fao.org/docrep/009/a0450e/a0450e00.htm. Viewed 12 May 2020.
- 550 11. Leung, B., Springborn, M. R., Turner, J. A., & Brockerhoff, E. G. (2014). Pathway-level
551 risk analysis: the net present value of an invasive species policy in the US. *Frontiers in
552 Ecology and the Environment*, 12(5), 273-279.
553
- 554 12. Hilbert, D. R., Roman, L. A., Koeser, A. K., Vogt, J., & van Doorn, N. S. (2019). Urban
555 tree mortality: a literature review. *Arboriculture & Urban Forestry*: 45 (5): 167-200., 45(5),
556 167-200.
- 557 13. Paap, T., Burgess, T. I., & Wingfield, M. J. (2017). Urban trees: bridge-heads for forest
558 pest invasions and sentinels for early detection. *Biological Invasions*, 19(12), 3515-3526.
559
- 560 14. Kovacs, K., Václavík, T., Haight, R. G., Pang, A., Cunniffe, N. J., Gilligan, C. A., &
561 Meentemeyer, R. K. (2011). Predicting the economic costs and property value losses
562 attributed to sudden oak death damage in California (2010–2020). *Journal of
563 Environmental Management*, 92(4), 1292-1302.
564
- 565 15. Koch, F. H., Ambrose, M. J., Yemshanov, D., Wiseman, P. E., & Cowett, F. D. (2018).
566 Modeling urban distributions of host trees for invasive forest insects in the eastern and
567 central USA: A three-step approach using field inventory data. *Forest Ecology and
568 Management*, 417, 222-236.
- 569 16. Potter, K. M., Escanferla, M. E., Jetton, R. M., & Man, G. (2019). Important insect and
570 disease threats to United States tree species and geographic patterns of their potential
571 impacts. *Forests*, 10(4), 304.

- 572 17. McGeoch, M. A., Genovesi, P., Bellingham, P. J., Costello, M. J., McGrannachan, C., &
573 Sheppard, A. (2016). Prioritizing species, pathways, and sites to achieve conservation
574 targets for biological invasion. *Biological Invasions*, 18(2), 299-314.
- 575 18. Sardain, A., Sardain, E., & Leung, B. (2019). Global forecasts of shipping traffic and
576 biological invasions to 2050. *Nature Sustainability*, 2(4), 274-282.
- 577
578 19. Haack, R. A., Hérard, F., Sun, J., & Turgeon, J. J. (2010). Managing invasive populations
579 of Asian longhorned beetle and citrus longhorned beetle: a worldwide
580 perspective. *Annual review of entomology*, 55.
- 581 20. Lovett, G. M., Weiss, M., Liebhold, A. M., Holmes, T. P., Leung, B., Lambert, K. F., ... &
582 Weldy, T. (2016). Nonnative forest insects and pathogens in the United States: Impacts
583 and policy options. *Ecological Applications*, 26(5), 1437-1455.
- 584
585 21. Knight, K. S., Brown, J. P., & Long, R. P. (2013). Factors affecting the survival of ash
586 (*Fraxinus* spp.) trees infested by emerald ash borer (*Agrilus planipennis*). *Biological*
587 *Invasions*, 15(2), 371-383.
- 588 22. Fei, S., Morin, R. S., Oswalt, C. M., & Liebhold, A. M. (2019). Biomass losses resulting
589 from insect and disease invasions in US forests. *Proceedings of the National Academy of*
590 *Sciences*, 116(35), 17371-17376.
- 591 23. Kinloch Jr, B. B. (2003). White pine blister rust in North America: past and
592 prognosis. *Phytopathology*, 93(8), 1044-1047.
- 593
594 24. Rizzo, D. M., & Garbelotto, M. (2003). Sudden oak death: endangering California and
595 Oregon forest ecosystems. *Frontiers in Ecology and the Environment*, 1(4), 197-204.
- 596
597 25. Coleman, T. W., Poloni, A. L., Chen, Y., Thu, P. Q., Li, Q., Sun, J., ... *et al.* (2019).
598 Hardwood injury and mortality associated with two shot hole borers, *Euwallacea* spp., in
599 the invaded region of southern California, USA, and the native region of Southeast
600 Asia. *Annals of Forest Science*, 76(3), 1-18.
- 601
602 26. Coutts, S. R., Helmstedt, K. J., & Bennett, J. R. (2018). Invasion lags: The stories we tell
603 ourselves and our inability to infer process from pattern. *Diversity and*
604 *Distributions*, 24(2), 244-251.
- 605
606 27. Hijmans, R. J., Phillips, S., Leathwick, J., Elith, J., & Hijmans, M. R. J. (2017). Package
607 'dismo'. *Circles*, 9(1), 1-68.
- 608 28. Fick, S. E., & Hijmans, R. J. (2017). WorldClim 2: new 1-km spatial resolution climate
609 surfaces for global land areas. *International Journal of Climatology*, 37(12), 4302-4315.
- 610 29. Homer, C.G., Dewitz, J.A., Yang, L., Jin, S., Danielson, P., Xian, G., ... *et al.* (2015.)
611 Completion of the 2011 National Land Cover Database for the conterminous United
612 States – representing a decade of land cover change information. *Photogrammetric*
613 *Engineering and Remote Sensing*, 81, 345–354.
- 614 30. Amodio, S., Aria, M., & D'Ambrosio, A. (2014). On concavity in nonlinear and
615 nonparametric regression models. *Statistica*, 74(1), 85-98.
- 616 31. Lambert, D. (1992). Zero-inflated Poisson regression, with an application to defects in
617 manufacturing. *Technometrics*, 34(1), 1-14.

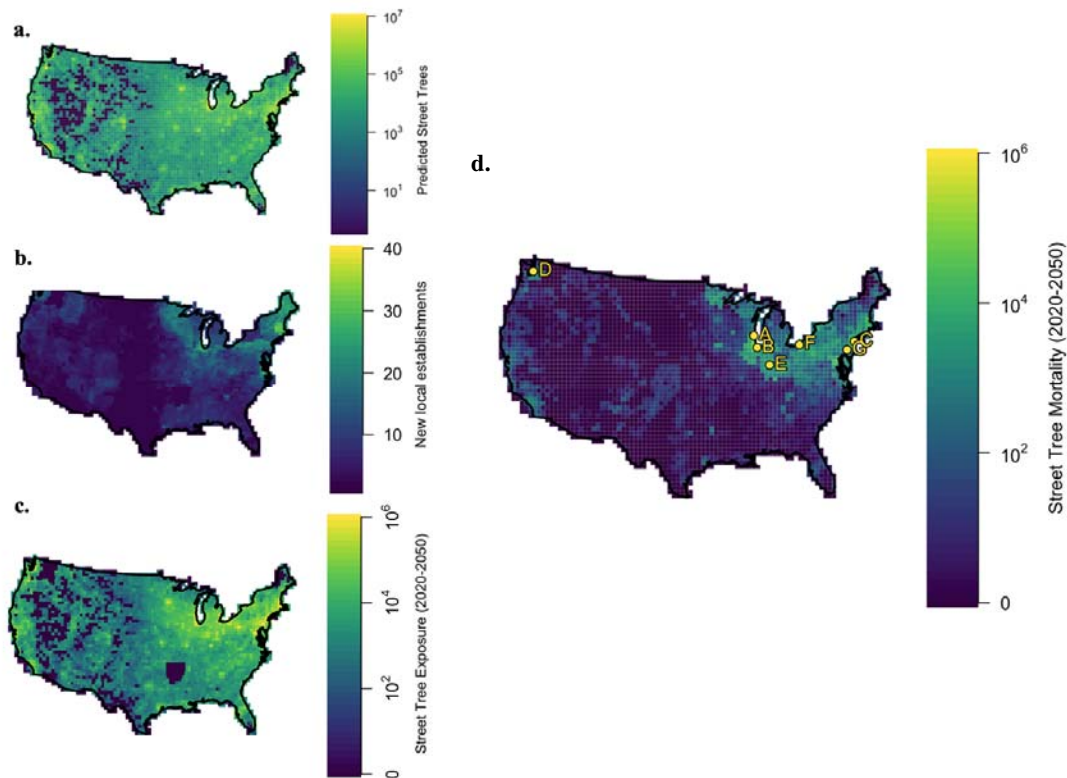
- 618 32. Wood, S. (2012). *mgcv: Mixed GAM Computation Vehicle with GCV/AIC/REML*
619 *smoothness estimation*. R package.
- 620 33. Elith, J., Leathwick, J. R., & Hastie, T. (2008). A working guide to boosted regression
621 trees. *Journal of Animal Ecology*, 77(4), 802-813.
- 622 34. Aukema, J. E., McCullough, D. G., Von Holle, B., Liebhold, A. M., Britton, K., & Frankel,
623 S. J. (2010). Historical accumulation of nonindigenous forest pests in the continental
624 United States. *BioScience*, 60(11), 886-897.
- 625 35. Carpenter, B., Gelman, A., Hoffman, M. D., Lee, D., Goodrich, B., Betancourt, M., ... *et al.*
626 (2017). Stan: A probabilistic programming language. *Journal of Statistical*
627 *Software*, 76(1).
- 628 36. Pugh, S.A. (2010). *Michigan's forest resources, 2008*. Research Note NRS-50. Newtown
629 Square, PA: U.S. Forest Service, Northern Research Station. 4 pp.
- 630
- 631 37. Hochberg, M.E., & Weis, A.E. (2001). Ecology: bagging the lag. *Nature* 409, 992–93.
- 632 38. Liebhold, A.M., & Tobin, P.C. (2008). Population ecology of insect invasions and their
633 management. *Annual Review of Entomology*, 53, 387–408.

634 **Figures and Tables**
635



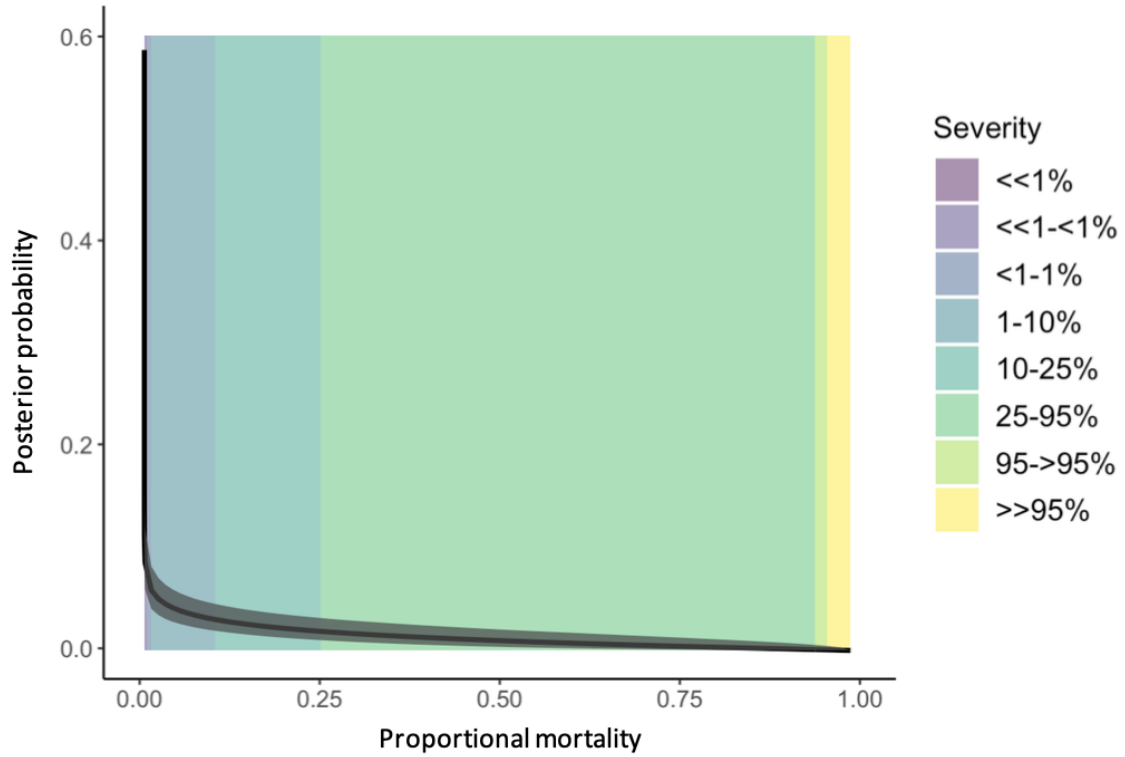
636
637
638

Figure 1. Fit of the genus-specific host tree models across all genera and size classes.



639
640
641
642
643
644
645
646
647

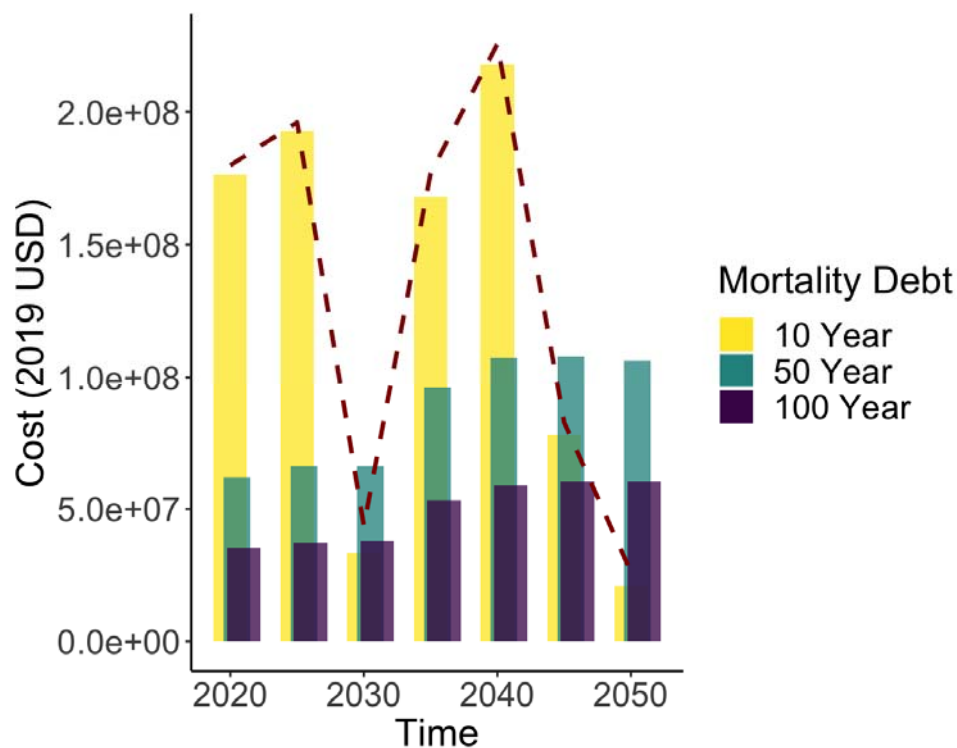
Figure 2. Model outputs for the first three subcomponent models, including **a.** predicted street tree abundance, **b.** predicted newly invaded sites of existing IAFIs, **c.** predicted street tree exposure levels (number of focal host tree + IAFI interactions) from 2020 to 2050, and finally **d.** Predicted total tree mortality from 2020 to 2050 in the most likely mortality debt scenario across space. The top seven most impacted cities or groups of nearby cities are shown in terms of total tree mortality 2020 to 2050 (A = Milwaukee, WI; B = Chicago/Aurora/Naperville/Arlington Heights, IL; C = New York, NY; D = Seattle, WA; E = Indianapolis, IN; F = Cleveland, OH; G = Philadelphia, PA).



648

649

650 **Figure 3.** Posterior distribution for the beta model of host mortality due to IAFIs within each
651 severity category. 95% Bayesian credible intervals are shown in grey, and the posterior median is
652 shown in black. Colored bins represent severity categories extended from [14].



653
654
655
656
657
658

Figure 4. Depiction of the influence of mortality debt on temporal cost patterns. Predicted costs 2020 to 2050 for the 10 year (yellow), 50 year (teal), and 100 year (purple) mortality debt scenarios with a 10 year initial invasion lag. The most likely scenario predictions are shown as a dashed red line. Costs are presented in 5-year increments in accordance with the timestep length within our spread model.

659 **Table 1.** Predicted annualized cost (in 2019 US dollars) and tree mortality across invasion
 660 scenarios from 2020 to 2050 across all 57 IAFI species. “Most likely” indicates the scenario with
 661 expert-elicited mortality debt durations by feeding guild, “Vary” scenarios hold all guilds but the
 662 focal guild constant at their most likely scenario, and “All” fix all three guilds at a given mortality
 663 debt duration. Mean mortality for most likely scenario = 2.3%, 1.38M trees, US\$ 30M annualized
 664 (US\$ 679M over the next 30 years).
 665

Mortality Debt Scenario	Annualized Cost (US\$ millions)		Tree Mortality (Millions)		Percent Mortality	
	lower 95% CI	upper 95% CI	lower 95% CI	upper 95% CI	lower 95% CI	upper 95% CI
Most likely	28.5	33.2	1.29	1.54	2.1%	2.5%
Vary Borers	10.1	32.1	0.45	1.45	0.7%	2.4%
Vary Defoliators	28.1	32.6	1.28	1.48	2.1%	2.4%
Vary Sap-feeders	28.5	32.5	1.30	1.47	2.1%	2.4%
All 10	27.8	30.4	1.27	1.39	2.1%	2.3%
All 50	18.5	22.3	0.84	1.00	1.4%	1.7%
All 100	9.77	13.5	0.44	0.60	0.7%	1.0%

666