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Do Species Distribution Models Predict Species Richness in Urban and Natural Green Space? A Case Study Using Amphibians

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Do species distribution models predict species richness in urban and natural green spaces? A case study using amphibians

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*Highlights

Research Highlights:

- Amphibian species richness maps significantly over-predicted species richness.

- Over-prediction may have partially been a result of undersampling during surveys.

- Over-prediction was likely due to poor model performance and undersampling.

- Despite over-prediction, models did project relative species richness well.

1   1. Introduction

2   Much of global population growth is occurring in urban areas (United Nations, 2004; Wu et

3   al. 2011) where over one-half of the United States population resides in urban areas (MacKun &

4   Wilson, 2011). Nevertheless, human land use patterns are dynamic and some locations within

5   urban areas are experiencing declining populations. Such declines can result in land

6   abandonment and provide an opportunity to replace developed habitat with green infrastructure.

7   It is well established that urbanization changes the biotic and abiotic properties of an ecosystem

8   and these impacts can reach far outside the urban area (Gaston, 2010). To reduce these effects,

9   there has been a movement to implement green infrastructure or incorporate green space to urban

10   areas. The benefits of green space in urban systems include increased psychological well-being,

11   recreational opportunities, and human health benefits (e.g., Barton & Pretty, 2010; Breuste &

12   Qureshi, 2011; Tzoulas et al., 2007; van den Berg, Hartig, & Staats, 2007). These benefits are

13   often predicated by ecosystem services and functions such green space in urban ecosystems

14   provide (e.g., water filtration and quality; Bolund & Hunhammar, 1999; Faulkner, 2004; Gaston,

15   Davies, & Edmondson, 2010). Whereas urban habitats may not act as smaller versions of

16   undeveloped patches of land, they may still provide ecological and human-oriented benefits such

17   as providing habitat connectivity, which helps to sustain regional biodiversity (Goddard, Dougill

18   & Benton, 2009; Irvine et al., 2010; Luck & Smallbone, 2010), or providing permeable surface

19   for stormwater infiltration, or water purification (Boyer & Polasky, 2004). Monitoring and

20   management for biodiversity has inherent value (Connery, 2009) and biodiversity conservation

21   within urban areas can help minimize extinction risk of some species and increase the value of

22   biota to humans as they more frequently encounter wildlife (Goddard et al., 2009). Toward this

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23   end, metrics are needed to measure the degree to which urban green spaces sustain biota and

24   subsequent biodiversity.

25   An ideal metric would use taxa that serve as indicators of overall biodiversity, provide an

26   ecosystem service, and are a critical link to the biotic community within the green space (i.e.,

27   provide ecosystem functions). Amphibians are often the most abundant, diverse group of

28   vertebrate organisms in forested and wetland systems, they serve as important food resources for

29   higher trophic levels, and in many systems are considered the top-predators (Burton & Likens,

30   1975; Davic & Welsh, 2004; Gibbons et al., 2006). Amphibians are also considered to be

31   indicators of environmental stress (DeGarady & Halbrook, 2006; Southerland et al., 2004; Welsh

32   & Droege, 2001; Welsh & Ollivier, 1998), but see Kerby et al. (2010), and are known to provide

33   a number of ecosystem functions in natural ecosystems (Davic & Welsh, 2004; Regester, Lips &

34   Whiles, 2006; Regester & Whiles, 2006; Whiles et al., 2006). Moreover, amphibians in urban

35   environments, like other biota, can enhance educational opportunities for human inhabitants

36   (Pickett et al., 2001). Because of their importance to ecosystems, ability to indicate

37   environmental stress, and education value, research involving amphibians in urban systems is

38   warranted (Hamer & McDonnell, 2008; McDonnell & Hahs, 2008; Pickett et al., 2001;

39   Smallbone, Luck & Wassens, 2011). Due to time and financial constraints associated with

40   conducting biotic surveys, modeling methods may provide assistance in understanding the value

41   of urban green space to this taxon.

42   Presence-only species distribution models (SDMs) are models that correlate species

43   distribution records to environmental data to predict areas of suitable habitat for taxa (see review

44   in Guisan & Zimmerman, 2000; Guisan & Thuiller, 2005; Elith et al., 2006). They are a group

45   of approaches for identifying species distributions of undersampled species, predicting impacts

46   of environmental change on distributions, and identifying areas of conservation importance

47   (Elith & Leathwick, 2009). In recent years, validations of models in various forms have been

48   increasing. For example, methods utilizing species occupancy or detection (Franklin et al., 2009;

49   Rota et al., 2011), independent and non-independent data validation (Araujo et al., 2005), and the

50   incorporation of field/survey data to inform or test model accuracy (Newbold et al., 2010; Pineda

51   & Lobo, 2009; Trotta-Moreu & Lobo, 2010) have been examined. However, studies simply

52   using field data to validate whether models are projecting species distribution correctly are rare.

53   Others have noted several limitations to SDMs including exclusion of biotic, geographical, or

54   physiological constraints on species distributions, use of museum records that may be widely

55   variable in both spatial and temporal quality, and issues relating to extrapolation of model

56   predictions (see review in Elith & Leathwick, 2009). These limitations may be exacerbated

57   when modeling within spatial extents that include urban environments, because species are

58   sampled less in urban areas as ecologists tend to focus collections or research on natural areas

59   (Gaston et al., 2010; Martin, Blossey & Ellis, 2012). Furthermore, lack of uniform sampling

60   across gradients of development presents a challenge to using SDMs in an urban landscape

61   because SDMs assume that biases in locality data (e.g., false absences) are not correlated with

62   environmental gradients used to build projected distributions (Bean, Stafford & Brashares, 2012;

63   Hijmans, 2012). In addition, error in the predictions of SDMs varies over large spatial scales

64   (extent and resolution) due to increased spatial heterogeneity (Osborne, Foody & Suarez-Seoane,

65   2007; Smulders et al., 2010; Zhang & Zhang, 2007), such as variation of environmental,

66   landscape, and habitat structure. This trend may be seen at smaller spatial scales (extent and

67   resolution) when using fine-scale data to build models (e.g., 30 m resolution) such as in urban

68   areas that have several classes of land use categories (e.g., habitat heterogeneity), as the

69   increased heterogeneity in urban areas within a smaller spatial scale could pose similar

70   prediction errors.

71   We tested whether species richness maps generated from SDMs can be used to prioritize areas

72   of high biodiversity value in urban and non-urban green space. We asked if SDMs built using

73   landscape variables associated with amphibian species richness could be used to project areas of

74   suitable habitat. We addressed this question by comparing modeled species richness maps

75   (based on accumulated individual SDMs) to field surveys across a number of urban and non-

76   urban green spaces. In addition, we investigated the landscape-level predictors of observed

77   amphibian species richness to determine what variables may be important to include or protect in

78   the creation, management, or conservation of urban and non-urban green space.

79

80   2. Materials and Methods

81   *2.1. Species Distribution Modeling using Maximum Entropy*

82   We developed species distribution models using Maxent version 3.3.3a (Phillips & Dudik,

83   2008) for 23 species of amphibians with current distributions within Hamilton County, Ohio,

84   U.S.A. Maxent is a software program that employs a machine learning method that is based on

85   the principle of maximum entropy to model species distributions using presence-only data

86   coupled with environmental data. Entropy is characterized by Shannon (1948) as "a measure of

87   how much „choice" is involved in the selection of an event" and is utilized in the framework of

88   maximum entropy to examine species geographic distributions (Phillips, Anderson & Schapire,

89   2006). The approach estimates habitat suitability based on an input set of environmental

90   variables encompassing the region where a species is known to occur based on locality records.

91   The program maximizes the entropy in the probability distribution of suitability across all areas

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92 of the distribution where empirical observations are lacking. For each species identified as

93 occurring in Hamilton County, OH, species presence data were obtained for the period of 1997–

94 2001from HerpNET (http://www.herpnet.org), Global Biodiversity Information Facility (Lane,

95 2003; GBIF; http://www.gbif.org), and personal collections of herpetologists (Appendix A). All

96 locality points were cross-referenced to each other and duplicate points were removed.

97 Furthermore, localities that fell outside the current species range (identified by county-level

98 distribution maps found in Lannoo, 2005) were not utilized to develop models. To maximize

99 model quality, each model was built using at least 20 point locations for each species (Wisz et

100 al., 2008).

101   We modeled the suitable habitat of each species across the National Hydrography Dataset

102 Plus (U.S. Geological Survey, 2005; NHDPlus; retrieved from http://www.horizon-

103 systems.com/nhdplus/) Region 05 Unit B watershed delineation. This delineation was necessary

104 to encompass the environmental variability of each species and provide predicted records from

105 across a larger geographic region adjacent to our primary study area. Additionally, amphibian

106 distributions can be restricted by geographic barriers (i.e., large rivers), and this region

107 encompassed a large portion of each species range, as well as the primary area of our field

108 surveys used to validate our models. This watershed delineation also encompasses a number of

109 similar adjacent urban areas (e.g., Lexington, KY, Indianapolis, IN, and Huntington, WV;

110 Appendix B) that could be examined in future studies.

111   We chose 11 initial environmental variables that others have shown to be important to

112 amphibians (e.g., Herrmann et al., 2005; Weyrauch & Grubb Jr, 2004) and had available data.

113 Geographic data layers were at a resolution of 30 m and variables extracted from these included

114 land cover, elevation, canopy cover, distance from stream, and slope/aspect. We created three

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115   layers using a moving window analysis to compute a ratio of cells classified as urban or water

116   within moving windows of 150, 300, and 500 m. Lastly, we calculated the number of cells

117   representing wetland habitat, as defined by the National Wetlands Inventory layer (U.S. Fish and

118   Wildlife Service, 2011; retrieved from http://www.fws.gov/wetlands/), within 300 m and 2 km

119   moving windows. We tested collinearity of the 11 layers by extracting environmental

120   information from 1000 randomly selected points (selected using the Random Point Generation

121   Tool within the Hawth"s Analysis Tool software for ArcGIS) within the watershed. A

122   correlation matrix was generated and correlations with $r \geq 0.70$ were considered highly

123   correlated. When pairs of variables exceeded this threshold ($r \geq 0.70$), we chose one variable

124   from the pair that we considered the most biologically relevant. Eight variables (listed below

125   with the data source) were chosen for inclusion in the final distribution model. Land cover

126   (Homer et al., 2007; retrieved from http://www.mrlc.gov/) was reclassified into one of seven

127   classes as follows (original land cover classes given in parenthesis): „forest" (41, 42, and 43),

128   water/wetlands (11, 90, and 95), „low intensity developed" (21 and 22), „medium intensity

129   developed" (23), „high intensity developed" (24), „natural non-forest" (31, 52, and 71), and

130   „agriculture" (81 and 82). Canopy cover values were taken from the National Land Cover

131   Database (NLCD; Homer et al., 2007; retrieved from http://www.mrlc.gov/). Elevation data

132   were derived from NHDPlus (National Hydrography Dataset; U.S. Geological Survey, 2005;

133   retrieved from www.horizon-systems.com/nhdplus/). The distance from stream (DSL) was

134   measured as distance (in meters) from the nearest drainage. Drainages were delineated from the

135   flow accumulation layer (downloaded from NHDPlus; retrieved from www.horizon-

136   systems.com/nhdplus/) as areas that drained 100 ha or more. The synthetic slope/aspect layer

137   (TASL; which represents aspect ranging from -1 [NE] to 1 [SW] and is weighted based on the

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138   steepness of the slope), was derived from NHDPlus elevation data following Pierce et al. (2005).

139   Finally, we selected three layers from the moving window analysis: Sum of urban, water and

140   wetland cells within a 300-m buffer. Urban and water cells were designated from our

141   reclassified NLCD database; whereas wetland cells were designated from the National Wetlands

142   Inventory layer (U.S. Fish and Wildlife Service, 2011; retrieved from

143   http://www.fws.gov/wetlands/). We chose the 300-m buffer because a 290-m buffer is

144   recommended for maintaining wetland and riparian habitat, thus this distance has been suggested

145   to be important for encompassing the core habitat requirement for many amphibian populations

146   (Semlitsch & Bodie, 2003). Moving window, DSL, and TASL layer calculations were

147   performed using ArcGIS (version 9.3; ESRI, Redlands, CA).

148   Maxent was run from the command line using the default settings with exception of

149   background points. A total of 620 target-group background data points representing localities of

150   amphibians in the designated watershed were used to develop an initial environmental envelope

151   to represent the range of environmental conditions within the modeled region. This method is

152   used to reduce the bias inherent in sample locality data (Phillips et al., 2009). The target-group

153   background approach uses background data (known as pseudo-absences) to develop the models.

154   The background points are chosen with the same biases as occurrence data and produce an

155   unbiased estimate of the geographic distribution of each species. The resulting background data

156   provide an equitable sample of the environmental conditions within the region modeled. Due to

157   the use of background points, we disabled the "addsamplestobackground" function. Further, we

158   disabled the "extrapolate," "Do MESS with analysis when projecting," and "Do clamping"

159   functions. Following methods in Milanovich et al. (2010), a multiple threshold approach was

160   used to designate a location as environmentally suitable for a species. Because Maxent produces

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161   a continuous probability of suitable habitat for each species, it is logistically unfeasible to present

162   each location as a probability of occupancy; therefore, we converted the continuous suitability

163   surface [0–1 from Maxent to presence/absence (1/0)] using four model output thresholds applied

164   by Maxent; fixed cumulative value 10 (FC10), minimum training presence (MTP), 10 percentile

165   training presence (10% TP), and maximum training sensitivity plus specificity (MTSPS). Next,

166   we generated four species richness maps based on the accumulated binary modeled distributions

167   of each species using the four Maxent thresholds. This four-threshold approach makes our

168   results comparable to other studies that provide predictions based on strict environmental

169   distributions of species (i.e., thresholds that maximize the agreement between observed and

170   predicted distributions; Cramer, 2003). This approach allows us to present model predictions

171   that relax the assumption of strict environmental control on species" distributions, and provides a

172   range of scenarios that could influence the predicted suitable habitat at the species-level and

173   comprehensive species richness.

174   We used null models to test the significance of each species distribution model (Raes & ter

175   Steege, 2007). We generated five null data sets, each with 1000 sets of sample points that were

176   randomly drawn without replacement from the pool of 620 background points. We generated a

177   null data set with the number of random points per distribution equal to 20, 45, 75, 150 and 250

178   data points, which represented a range of presence points available to model each species.

179   Maxent was used to calculate the area under the curve (AUC) for the 1000 null data sets to create

180   an AUC frequency distribution. The calculated AUC for each species model was compared to

181   the 95th percentile AUC value of the null frequency distribution created from the representative

182   number of sample points (20, 45, 75, 150, or 250). A species model performs better than random

183   (e.g., null model) and is considered significant if the calculated AUC is greater than the

184   corresponding 95th percentile AUC of the null-distribution (Raes & ter Steege, 2007).

185   *2.2. Examining differences in amphibian species richness*

186   *2.2.1. Amphibian surveys*

187   To evaluate the degree to which SDMs accurately predict amphibian richness, we conducted

188   amphibian surveys at 20 sites across the Cincinnati, OH metropolitan area following methods

189   proposed in Shaffer et al. (1994). Each site consisted of identified green space, spanned a range

190   of sizes (3–1758 ha; Appendix C), and were spread across a gradient of urbanization (e.g., within

191   and outside the designated Cincinnati, OH metropolitan area; Fig. 1; Appendix B). Surveys were

192   conducted March – June 2011 and each site was sampled three times with a 3–5 week period

193   between repeat samples. To standardize our search effort within each site, nine plots were

194   selected and surveyed at each site. We attempted to place three plots in each of three habitat

195   types associated with amphibian richness: Wetlands/ponds, streams, and terrestrial (forest or

196   non-forest natural [e.g., grassland/prairie]) habitat. In the event that three each of

197   wetlands/ponds or streams were not available, we increased the number of terrestrial plots to

198   reach nine total plots for each site. If more than three of an aquatic habitat type were available

199   (e.g., wetlands/ponds or streams) we sampled plots with the most suitable amphibian habitat (i.e.,

200   forested wetlands/ponds or streams). We employed three survey techniques at each site: area

201   sensitive dip-net surveys for pond and/or ephemeral wetland habitats, area sensitive terrestrial

202   (searching cover objects) surveys, and leaf litter bags and dip-netting area sensitive surveys for

203   headwater streams following Chalmers & Droege (2002) and Waldron et al. (2003). At each

204   wetland/pond, we dip-netted several 1 m^2 areas adjacent to the bank using a 40.6 X 40.6 cm (3.2

205   mm mesh size) net. Plots were placed every 10 m until the entire perimeter of the body of water

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206   was sampled. For terrestrial plots, we surveyed five, 5 m^2 sub-plots at each site. Plots were

207   chosen by placing a single 5 m^2 plot within the center of each habitat type occurring on the site

208   (forest or non-forest natural [prairie or grassland habitats]) and an additional 5 m^2 plot in each

209   cardinal direction 25 m from the center plot. Within each sub-plot, all cover objects and existing

210   leaf litter were searched. Within each stream, five 1 m^2 plots were created every 10 m starting

211   approximately 20 m from the confluence of a larger stream, road, or trail and continuing

212   upstream. Within each plot, we placed one leaf litter bag (55 x 25 cm made with 1.3 cm^2 mesh)

213   filled with deciduous leaf litter from the surrounding forest in a wetted portion of the stream

214   channel with a large rock on top to prevent dislodging. We followed the methods of

215   Nowakowski & Maerz (2009) and Peterman, Truslow, & Samuel (2008) to remove contents

216   from each bag; we thoroughly dip-netted the 1 m^2 plot by dislodging substrate parallel to the

217   stream bank across the width of the stream. In addition to our structured surveys, any animals

218   encountered (seen or heard) while travelling between habitat types or plots within a site were

219   noted. All animals captured or heard (frog calls) were identified to species.

220   *2.2.2. Validation of amphibian surveys and richness comparisons*

221   We examined how well predicted species richness matched observed species richness by

222   comparing the maximum species richness value from each site between observed and predicted

223   (from each Maxent threshold) values derived from our maps. We compared differences between

224   species richness within "urban" and "non-urban" sites by categorizing a site as "urban" if $\geq 50\%$

225   of the adjacent landscape within a 2-km buffer of the centroid of the site had urban cells (NLCD

226   categories $21 - 24$) based on our NLCD 2001 reclassified maps. To test for differences between

227   predicted species richness from each model threshold, we used paired *t*-tests to compare

228   observed and predicted (from each threshold) species richness values from each site surveyed.

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229   To examine if we adequately sampled species richness, we estimated species richness using

230   EstimateS version 8.2 (Colwell, 2009) for six sites that had been surveyed on at least 10

231   occasions. Additional survey data were derived from reports summarized in Davis, Krusling,

232   and Ferner (1998). These reports summarized data from county-park-level surveys that took

233   place between 1988 and 1995, surveyed a similar level of habitat heterogeneity and utilized

234   similar survey methods to our study, and lasted approximately one year. We chose to make

235   comparisons between species richness at this group of sites using the Chao 2 estimator because

236   this algorithm produces species accumulation curves that approach maximum values with few

237   samples (i.e., the estimator was the least sensitive to undersampling; Colwell & Coddington,

238   1994). The Chao 2 algorithm inflates the observed species richness by a factor derived from the

239   number of species observed only once or twice within a total sample. The estimator is calculated

240   as $S_{Chao2} = S_{obs} + Q^2_1 / 2Q_2$, where S_{obs} is the observed species richness and Q_1 and Q_2 are the

241   number of species detected only once or twice per site, respectively. Thus, this estimator

242   accounts for the fact that species differ in detectability and uses the relative frequencies of

243   species that are rarely detected to estimate the number of taxa that are present but not detected

244   (Chao, 1987; Colwell & Coddington, 1994). For each additional site we calculated coverage

245   (number of species observed/number of species estimated X 100), exclusive species (percentage

246   of species only observed in a given site or category), and completeness of a sample (species

247   observed as a percentage of the total number of species expected in the site; Gardner et al.,

248   2007). This approach estimated the amphibian species richness at sites visited ≥ 10 times and

249   provided an estimate of the number of sampling periods needed to accurately capture the

250   estimated richness value. These values were then compared to the observed amphibian species

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251 richness values derived from the current surveys and the surveys of Davis, Krusling, and Ferner

252 (1998).

253   *2.2.3. Identifying* f*actors that influence observed species richness*

254   We used a general linear model and an information theoretic approach to examine which

255 environmental factors were most predictive of observed amphibian species richness. This model

256 is specific to existing green spaces, unlike the species distribution models, which provide a more

257 general estimate of species richness patterns. As a result, the environmental variables used in

258 this analysis are similar to those in the distribution model; however, some differences do exist.

259 Observed species richness was the dependent variable. Independent variables included the

260 percent of urban land within a 2-km buffer of the centroid of each site (square root transformed),

261 the number of wetlands within a 2-km buffer of the centroid of each site (log transformed),

262 percent forest within each site (square root transformed), and percent non-forest natural habitat

263 (agriculture and non-forested natural habitat) within each site (square root transformed) as

264 continuous predictor variables. Using the same variables as above, we evaluated the importance

265 of landscape variables on observed species richness using Akaike Information Criterion

266 (Burnham & Anderson, 2002). These variables were derived from our reclassified NLCD 2001

267 land-cover layer and a National Wetlands Inventory layer (U.S. Fish and Wildlife Service, 2011;

268 http://www.fws.gov/wetlands/) and were chosen out of an initial set of 11 independent variables

269 that we thought would have an influence on amphibian species richness (Table 3). We generated

270 a correlation matrix of all 11 variables and chose four variables that were not correlated with

271 each other ($r \leq 0.70$). Variables were transformed to meet normality assumptions (Sokal &

272 Rohlf, 1995).

12

273   Lastly, to determine the relationship between our observed species richness values and the

274   mean predicted species richness values across the urban gradient, we conducted an analysis of

275   covariance. Maxent predictions (predicted species richness) and field observations (observed

276   species richness) were factors in the analysis, percent urbanization within 2-km of each site

277   (square root transformed) was our covariate, and observed species richness and mean predicted

278   species richness values were used as our dependent variables. The significance of the interaction

279   term was evaluated to determine if the relationship between observed and predicted species

280   richness estimates differed across the urban treatments.

281

282   3. Results

283   The mean AUC for amphibian distribution models based on landscape variables was 0.83

284   (range = 0.70 – 0.96; median = 0.83; Appendix A) and the AUC for each species was better than

285   random (i.e., model AUC values exceeded the 95th percentile of the null AUC distributions).

286   Observed and predicted species richness varied across each site (Appendix C). Mean observed

287   species richness was triple in non-urban sites compared to the value in urban sites (mean species

288   richness values; overall = 3.2, urban = 1.9, non-urban = 6.2). There were significant differences

289   between observed and predicted species richness for each threshold (Table 1; all t-tests $P \leq$

290   0.001). Overall, mean differences between observed and predicted species richness within each

291   threshold ranged from 253 to 558% (mean range = 8 to 18 species). These differences increased

292   when "non-urban" sites were isolated from "urban" sites (mean range = 343 to 619%; mean

293   range = 8 to 17 species) and decreased when sites sampled greater than 10 times were isolated

294   from sites sampled three times (mean range = 14 to 98%; mean range = 2 to 11 species; see

295   "Additional" column in Table 1).

13

296   The MTSPS threshold, our most conservative threshold, had the lowest levels of over-

297   prediction between observed and predicted species richness in both urban and non-urban sites;

298   whereas the MTP threshold had the greatest levels of over-prediction. The species accumulation

299   curves (Fig. 2) for all green spaces sampled > 3 times, except Farbach-Werner Nature Preserve,

300   suggest these sites had been (more or less) exhaustively sampled (i.e., the species accumulation

301   curve reaches a plateau). There was no site for which the final estimate of species richness from

302   the species accumulation curves was achieved by the third sample (based on, for example, Chao

303   2 richness estimator; Fig. 2; Table 2).

304   Observed amphibian species richness varied depending on the number of wetlands and

305   percent of urbanization within a 2-km buffer around the centroid of each site (Tables 3 and 4).

306   Percent of forest or non-forest natural habitat within each site did not predict observed species

307   richness (Table 3). The best model (based on lowest AICc and highest w_i) for observed species

308   richness also includes number of wetlands and percent of urbanization within a 2-km buffer

309   around the centroid of each site (Table 4). Percent of forest or non-forest natural habitat within

310   each site did not predict observed species richness (Table 3). Despite the significant

311   discrepancies in absolute richness values between the observed and predicted species richness

312   values, there was a significant trend of decreasing richness with increasing urban habitat ($P <$

313   0.001 for both observed and mean predicted richness). Further, the slopes of these two trends

314   did not differ ($P = 0.913$; Appendix D) between modeled and observed values and the

315   relationships between observed and predicted species richness was significantly correlated to the

316   level of adjacent urbanization (Appendix D).

317

318   4. Discussion and Conclusions

319 The use of SDMs built with high-resolution landscape variables and real-time amphibian

320 surveys allows us to assess whether this technique is a valid method to identify areas of higher

321 and lower species richness across urban and non-urban green space. Our approach of modeling

322 followed by field surveys revealed differences between observed and predicted species richness

323 across urban and non-urban green space (Table 1). These differences can be interpreted as (1)

324 sites were undersampled, thus our observed measure of species richness could be inaccurate, or

325 (2) the SDMs are over representing actual species distributions – which accumulate to unrealistic

326 predictions of species richness across the landscape. Our results suggest the differences are

327 likely due to both factors. First, species accumulation curves derived from the Chao 2 estimator

328 based on six sites that were extensively surveyed (i.e., ≥ 10 sampling events over approximately

329 one year), indicated three sampling periods was not a sufficient effort to capture species richness.

330 In each of these six sites, species accumulation curves did not plateau until between six to 10

331 sampling periods (Fig. 2). We conducted an intensive single-season survey that reasonably

332 mimics the effort many agencies would implement to understand factors driving species

333 distributions of amphibians. We know from studies with a more intensive sampling effort that

334 this is not sufficient to capture all species, because often species are unavailable for capture for

335 entire breeding years (e.g., Barrett & Guyer, 2008).

336 However, undersampling alone did not explain all of the differences between observed and

337 predicted species richness. Chao 2 species richness estimates showed SDMs still over-predicted

338 species richness even in sites considered to be adequately sampled (Table 2). For that subset of

339 sites, we have reasonable confidence (empirically demonstrated via richness estimator) that all

340 differences between sampled and predicted species richness are not due to undersampling alone.

341 This suggests the differences are at least partially a consequence of SDM predictions.

342   We propose the SDM influence on over-prediction is the result of the inherent nature of

343   SDMs. By solely using environmental variables to construct predictions of a species" suitable

344   habitat, SDMs fail to incorporate biological or geographic influence on species distributions

345   (e.g., Guisan & Thuiller, 2005; Heikkinen et al., 2006; Luoto et al., 2005), also referred to as

346   realized versus fundamental niche (Rodda, Jarnevich & Reed, 2011; Soberón & Peterson, 2005).

347   This in turn can lead to an overestimation of species suitable habitat, as only areas of suitable

348   habitat, not realized distribution, are projected. This overestimation may increase when adding

349   multiple individual models to create species richness maps (Hortal & Lobo, 2006; Pineda &

350   Lobo, 2009; Thullen, Sartoris & Walton, 2002; Vasconcelos, Rodriguez & Hawkins, 2011), as

351   was the case in this study. Furthermore, although the use of target-group background data is

352   suggested to reduce sampling bias (Phillips et al., 2009), use of this approach could be a possible

353   source of overestimation. Models developed utilizing replicate samples and random background

354   data could result in a greater or lesser over-estimation of species richness values. Further

355   investigation into the influence of various modeling approaches to overestimation of species

356   richness is warranted.

357   Using multiple thresholds to develop a range of scenario"s of species predicted suitable

358   habitat was a definite strength of this study (Fig. 1; Appendices B and C), and the differences

359   between predicted species richness for different model thresholds should not be ignored. The

360   importance of presenting a range of model thresholds has been supported in other studies (Liu et

361   al., 2005; Milanovich et al., 2010; Pineda & Lobo, 2009; Thullen et al., 2002). We suggest

362   presenting a range of thresholds allowed the geographic range of suitable habitat for each species

363   to be both larger and smaller than realized ranges. This may account for some of the geographic

364   and biotic influences on species ranges; however, as our results indicate, threshold adjustment

365   may be a necessary, but not entirely sufficient means for generating more precise SDMs.

366   Despite the over-prediction of our models and differences between model thresholds, we

367   argue this technique is a useful tool for management of green space by identifying areas suitable

368   for amphibians in urban ecosystems. Specifically, although the model richness estimate was

369   high, there was no difference between the trend in observed richness and modeled richness

370   across the urban treatments. Species distribution models are increasingly used to identify

371   potential areas for conservation and management of biodiversity (Lawler, Wiersma &

372   Huettmann, 2011; Loiselle et al., 2003). In this study, SDMs identified areas of higher

373   suitability, as green space with higher observed species richness generally had higher predicted

374   species richness (Appendix B and C). For example, the Cincinnati Nature Center and

375   Richardson Nature Preserve (the non-urban sites with the highest observed species richness) and

376   Farbach-Werner Nature Preserve and Triple Creek Park (the urban sites with the highest

377   observed species richness) each had some of the highest predicted species richness within each

378   threshold (Appendix C). One commonality between these sites is each site had a high number of

379   adjacent wetlands (within 2-km of the centroid), and therefore, a high number of aquatic habitats

380   surveyed (Appendix C). With respect to management of existing or development of new urban

381   green spaces, we found the number of wetlands adjacent to each site was a significant predictor

382   of observed amphibian richness (Tables 3 and 4). Our predictive models support the importance

383   of wetlands in urban green space to amphibian species richness, as the two urban sites with

384   surveyed wetlands (Farbach-Werner N.P. and Triple Creek Park) had the highest predicted

385   species richness of all urban sites. This highlights the importance of prioritizing the

386   maintenance, restoration, or mitigation of wetlands in urban green space. Furthermore, this is a

387   testament to the usefulness of SDMs for identifying areas of potential conservation importance,

17

388   but emphasizes that models must be constructed with biologically relevant variables that

389   facilitate presence of a particular species or taxon (e.g., wetlands or forest for amphibians).

390   In conclusion, we tested whether species distribution models could be utilized to identify

391   areas of higher amphibian species richness in urban ecosystems. If one"s goal is to make rapid

392   decisions concerning the management or creation of green space or infrastructure based on

393   conserving areas of higher biodiversity, this method provides relative measures of richness. Our

394   results indicate substantial effort is needed to sufficiently survey and identify areas of suitable

395   habitat for amphibians in urban and non-urban green spaces. Field surveys often require

396   expertise, time, energy, and funds; whereas SDMs can be developed using existing, readily

397   available data (e.g., museum records) and can be conducted using a variety of freely available

398   programs (Elith et al., 2006). We offer two suggestions on how to improve SDM use if one"s

399   goal is to identify areas important to high biodiversity. First, the development of species-specific

400   models, particularly in cases where there may be a desire to increase the abundance of a

401   particular species of concern, could enhance predictive ability. These models could utilize key

402   (spatially explicit) variables important to the biology of a particular species. Second, we

403   recommend using SDM approaches that incorporate an estimation of probability of occurrence

404   and species detectability, where such data are available for species of interest. This method has

405   been successfully utilized and detailed in more recent studies (e.g., Franklin et al., 2009;

406   Newbold et al., 2010; Rota et al., 2011). Unfortunately, our data did not permit such analyses.

407   We suggest incorporating one or both of these approaches to strengthen the accuracy of SDMs in

408   predicting species richness.

References

Araujo, M. B., Pearson, R. G., Thuiller, W. & Erhard, M. (2005). Validation of species-climate impact models under climate change. *Global Change Biology,* 11, 1504-1513.

Barrett, K. & Guyer, C. (2008). Differential responses of amphibians and reptiles in riparian and stream habitats to land use disturbances in western Georgia, USA. *Biological Conservation,* 141, 2290-2300.

Barton, J. & Pretty, J. (2010). Urban ecology and human health and wellbeing. In K. J. Gaston (Ed.). *Urban ecology* (pp. 202-229). New York, NY - Cambridge University Press.

Bean, W. T., Stafford, R. & Brashares, J. S. (2012). The effects of small sample size and sample bias on threshold selection and accuracy assessment of species distribution models. *Ecography,* 35, 250-258.

Bolund, P. & Hunhammar, S. (1999). Ecosystem services in urban areas. *Ecological Economics,* 29, 293-301.

Boyer, T. & Polasky, S. (2004). Valuing urban wetlands: a review of non-market valuation studies. *Wetlands,* 24, 744-755.

Breuste, J. & Qureshi, S. (2011). Urban sustainability, urban ecology and the Society for Urban Ecology (SURE). *Urban Ecosystems,* 14, 313-317.

Burnham, K. P. & Anderson, D. R. (Eds.). (2002). *Model selection and multimodel inference: a practical information-theoretic approach.* New York, NY - Springer-Verlag Press.

Burton, T. M. & Likens, G. E. (1975). Energy flow and nutrient cycling in salamander populations in the Hubbard Brook Experimantal Forest, New Hampshire. *Ecology,* 56, 1068-1080.

Chalmers, R. J. & Droege, S. (2002). Leaf litter bags as an index to populations of northern two-lined salamanders (*Eurycea bislineata*). *Wildlife Society Bulletin,* 30, 71-74.

Chao, A. (1987). Estimating the population size for capture-recapture data with unequal catchability. *Biometrics,* 43, 783-791.

Colwell, R. K. (2009). *EstimateS: statistical estimation of species richness and shared species from samples, Version 8.2.0.* Accessed online December 2011. http://viceroy.eeb.uconn.edu/estimates/.

Colwell, R. K. & Coddington, J. A. (1994). Estimating terrestrial biodiversity through extrapolation. *Philosophical Transactions of the Royal Society of London B Biological Sciences,* 345, 101-118.

Connery, K. (2009). Biodiversity and urban design: seeking an integrated solution. *Journal of Green Building,* 4, 23-38.

Cramer, J. S. (Ed.) (2003). *Logit models: from economics and other fields.* Oxford - Cambridge University Press.

Davic, R. D. & Welsh, H. H. J. (2004). On the ecological roles of salamanders. *Annual Review of Ecology Evolution and Systematics,* 35, 405-434.

Davis, J. G., Krusling, P. J. & Ferner, J. W. (1998). Status of amphibian populations in Hamiltion County, Ohio. In M. J. Lannoo (Ed.). *Status and conservation of midwestern amphibians* (pp. 155-165). Ames - University of Iowa Press.

DeGarady, C. J. & Halbrook, R. S. (2006). Using anurans as bioindicators of PCB contaminated streams. *Journal of Herpetology,* 40, 127-130.

Elith, J., Graham, C. H., Anderson, R. P., Dudik, M., Ferrier, S., Guisan, A., Hijmans, R. J., Huettmann, F., Leathwick, J. R., Lehmann, A., Li, J., Lohmann, L. G., Loiselle, B. A., Manion, G., Moritz, C., Nakamura, M., Nakazawa, Y., Overton, J. M., Peterson, A. T., Phillips, S. J., Richardson, K., Scachetti-Pereira, R., Schapire, R. E., Soberon, J., Williams, S., Wisz, M. S. & Zimmermann, N. E. (2006). Novel methods improve prediction of species' distributions from occurrence data. *Ecography,* 29, 129-151.

Elith, J. & Leathwick, J. R. (2009). Species distribution models: ecological explanation and prediction across space and time. *Annual Review of Ecology Evolution and Systematics,* 40, 677-697.

Faulkner, S. (2004). Urbanization impacts on the structure and function of forested wetlands. *Urban Ecosystems,* 7, 89-106.

Fish and Wildlife Service, U. S. (2011). *National Wetlands Inventory website.* Accessed December 2011. http://www.fws.gov/wetlands/.

Franklin, J., Wejnert, K. E., Hathaway, S. A., Rochester, C. J. & Fisher, R. N. (2009). Effect of species rarity on the accuracy of species distribution models for reptiles and amphibians in southern California. *Diversity and Distributions,* 15, 167-177.

Gardner, T. A., Ribeiro-Junior, M. A., Barlow, J., Cristina, T., Avila-Pires, S., Hoogmoed, M. S. & Peres, C. A. (2007). The value of primary, secondary, and plantation forests for a neotropical herpetofauna. *Conservation Biology,* 21, 775-787.

Gaston, K. J. (2010). Urban ecology. In K. J. Gaston (Ed.). *Urban ecology* (pp. 1-9). New York - Cambridge University Press.

Gaston, K. J., Davies, Z. G. & Edmondson, J. L. (2010). Urban environments and ecosystem functions. In K. J. Gaston (Ed.). *Urban ecology* (pp. 35-52). New York - Cambridge University Press.

Geological Survey, U. S. (2005). *The National Hydrography Dataset Plus - NHDPlus.* Accessed online December 2011. http://www.horizon-systems.com/nhdplus/.

Gibbons, J. W., Winne, C. T., Scott, D. E., Willson, J. D., Glaudas, X., Andrews, K. M., Todd, B. D., Fedewa, L. A., Wilkinson, L., Tsaliagos, R. N., Harper, S. J., Greene, J. L., Tuberville, T. D., Metts, B. S., Dorcast, M. E., Nestor, J. P., Young, C. A., Akre, T., Reed, R. N., Buhlmann, K. A., Norman, J., Croshaw, D. A., Hagen, C. & Rothermel, B. B. (2006). Remarkable amphibian biomass and abundance in an isolated wetland: implications for wetland conservation. *Conservation Biology,* 20, 1457-1465.

Goddard, M. A., Dougill, A. J. & Benton, T. G. (2009). Scaling up from gardens: biodiversity conservation in urban environments. *Trends in Ecology & Evolution,* 25, 90-98.

Guisan, A. & Thuiller, W. (2005). Predicting species distribution: offering more than simple habitat models. *Ecology Letters* 8, 993-1009.

Guisan, A. & Zimmermann, N. E. (2000). Predictive habitat distribution models in ecology. *Ecological Modelling,* 135, 147-186.

Hamer, A. J. & McDonnell, M. J. (2008). Amphibian ecology and conservation in the urbanising world: a review. *Biological Conservation,* 141, 2432-2449.

Heikkinen, R. K., Luoto, M., Araujo, M. B., Virkkala, R., Thuiller, W. & Sykes, M. T. (2006). Methods and uncertainties in bioclimatic envelope modelling under climate change. *Progress in Physical Geography,* 30, 751-777.

HerpNET. (2011). HerpNET. Accessed online December 2011. http://www.herpnet.org.

Herrmann, H. L., Babbitt, K. J., Baber, M. J. & Congalton, R. G. (2005). Effects of landscape characteristics on amphibian distribution in a forest-dominated landscape. *Biological Conservation,* 123, 139-149.

Hijmans, R. J. (2012). Cross-validation of species distribution models: removing spatial sorting bias and calibration with a null model. *Ecology,* 93, 679-688.

Homer, C., Dewitz, J., Fry, J., Coan, M., Hossain, N., Larson, C., Herold, N., McKerrow, A., VanDriel, J. N. & Wickham, J. (2007). Completion of the 2001 National Land Cover Database for the Conterminous United States. *Photogrammetric Engineering and Remote Sensing,* 73, 858-864.

Hortal, J. & Lobo, J. M. (2006). Towards a sysecological framework for systematic conservation planning. *Biodiversity Informatics,* 3, 16-45.

Irvine, K. N., Fuller, R. A., Devine-Wright, P., Tratalos, J., Payne, S. R., Warren, P. H., Lomas, K. J. & Gaston, K. J. (2010). Ecological and psychological value of urban green space. In M. Jenks & C. Jones Eds.). *Dimensions of the sustainable City* (pp. 296). New York - Springer Verlag

Kerby, J. L., Richards-Hrdlicka, K. L., Storfer, A. & Skelly, D. K. (2010). An examination of amphibian sensitivity to environmental contaminants: are amphibians poor canaries? *Ecology Letters,* 13, 60-67.

Lane, M. A. (2003). The global biodiversity informaiton facility. *Bulletin of the American Society for Information Science and Technology,* 30, 22-24.

Lannoo, M. J. (Ed.) (2005). *Amphibian declines: the conservation status of United States species.* Berkely, CA - University of California Press.

Lawler, J. J., Wiersma, Y. F. & Huettmann, F. (2011). Using species distribution models for conservation planning and ecological forecasting. In C. A. Drew, Y. F. Wiersma & F. Huettmann Eds.). *Predictive species and habitat modeling in landscape ecology: concepts and applications* (pp. 271-290). New York - Springer.

Liu, C. R., Berry, P. M., Dawson, T. P. & Pearson, R. G. (2005). Selecting thresholds of occurrence in the prediction of species distributions. *Ecography,* 28, 385-393.

Loiselle, B. A., Howell, C. A., Graham, C. H., Goerck, J. M., Brooks, T., Smith, K. G. & Williams, P. H. (2003). Avoiding pitfalls of using species distribution models in conservation planning. *Conservation Biology,* 17, 1591-1600.

Luck, G. W. & Smallbone, L. T. (2010). Species diversity and urbanisation: patterns, drivers and implications. In K. J. Gaston (Ed.). *Urban ecology* (pp. 88-119). New York - Cambridge University Press.

Luoto, M., Pöyry, J., Heikkinen, R. K. & Saarinen, K. (2005). Uncertainty of bioclimate envelope models based on the geographical distribution of species. *Global Ecology and Biogeography,* 14, 575-584.

MacKun, P. & Wilson, S. (2011). *Population Distribution and Change: 2000 to 2010.* U.S. Census Bureau. Accessed online December 2011. www.census.gov.

Martin, L. J., Blossey, B. & Ellis, E. (2012). Mapping where ecologists work: biases in the global distribution of terrestrial ecological observations. *Frontiers in Ecology and the Environment,* 10, 195-201.

McDonnell, M. J. & Hahs, A. K. (2008). The use of gradient analysis studies in advancing our understanding of the ecology of urbanizing landscapes: current status and future directions. *Landscape Ecology,* 23, 1143-1155.

Milanovich, J. R., Peterman, W. E., Nibbelink, N. P. & Maerz, J. C. (2010). Projected loss of a salamander diversity hotspot as a consequence of projected global climate change. *PLoS ONE,* 5, e12189. doi:10.1371/journal.pone.0012189.

Nations, U. (2004). The United Nations on world population in 2300. *Population and Development Review,* 30, 181-187.

Newbold, T., Reader, T., El-Gabbas, A., Berg, W., Shohdi, W. M., Zalat, S., El Din, S. B. & Gilbert, F. (2010). Testing the accuracy of species distribution models using species records from a new field survey. *Oikos,* 119, 1326-1334.

Nowakowski, A. J. & Maerz, J. C. (2009). Estimation of larval stream salamander densities in three proximate streams in the Georgia Piedmont. *Journal of Herpetology,* 43, 503-509.

Osborne, P. E., Foody, G. M. & Suarez-Seoane, S. (2007). Non-stationarity and local approaches to modelling the distributions of wildlife. *Diversity and Distributions,* 13, 313-323.

Peterman, W. E., Truslow, W. E. & Samuel, C. (2008). Density estimation of larval *Eurycea wilderae*: a comparison of mark-recapture and depletion sampling. *Herpetological Review,* 39, 438-442.

Phillips, S. J., Anderson, R. P. & Schapire, R. E. (2006). Maximum entropy modeling of species geographic distributions. *Ecological Modelling,* 190, 231-259.

Phillips, S. J. & Dudik, M. (2008). Modeling of species distributions with Maxent: new extensions and a comprehensive evaluation. *Ecography,* 31, 161-175.

Phillips, S. J., Dudik, M., Elith, J., Graham, C. H., Lehmann, A., Leathwick, J. & Ferrier, S. (2009). Sample selection bias and presence-only distribution models: implications for background and pseudo-absence data. *Ecological Applications,* 19, 181-197.

Pickett, S. T. A., Cadenasso, M. L., Grove, J. M., Nilon, C. H., Pouyat, R. V., Zipperer, W. C. & Costanza, R. (2001). Urban ecological systems: Linking terrestrial ecological, physical, and socioeconomic components of metropolitan areas. *Annual Review of Ecology and Systematics,* 32, 127-157.

Pierce, K. B., Lookingbill, T. & Urban, D. (2005). A simple method for estimating potential relative radiation (PRR) for landscape-scale vegetation analysis. *Landscape Ecology,* 20, 137-147.

Pineda, E. & Lobo, J. M. (2009). Assessing the accuracy of species distribution models to predict amphibian species richness patterns. *Journal of Animal Ecology,* 78, 182-190.

Raes, N. & ter Steege, H. (2007). A null-model for significance testing of presence-only species distribution models. *Ecography,* 30, 727-736.

Regester, K. J., Lips, K. R. & Whiles, M. R. (2006). Energy flow and subsidies associated with the complex life cycle of ambystomatid salamanders in ponds and adjacent forest in southern Illinois. *Oecologia,* 147, 303-314.

Regester, K. J. & Whiles, M. R. (2006). Decomposition rates of salamander (*Ambystoma maculatum*) life stages and associated energy and nutrient fluxes in ponds and adjacent forest in in Southern Illinois. *Copeia,* 640-649.

Rodda, G. H., Jarnevich, C. S. & Reed, R. N. (2011). Challenges in identifying sites climatically matched to the native ranges of animal invaders. *PLoS ONE,* 6, e14670. doi:10.1371/journal.pone.0014670.

Rota, C. T., Fletcher, R. J., Jr. , Evans, J. M. & Hutto, R. L. (2011). Does accounting for imperfect detection improve species distribution models? *Ecography,* doi: 10.1111/j.1600-0587.2010.06433.x.

Semlitsch, R. D. & Bodie, J. R. (2003). Biological criteria for buffer zones around wetlands and riparian habitats for amphibians and reptiles. *Conservation Biology,* 17, 1219-1228.

Shaffer, H. B., Alford, R. A., Woodward, B. D., Richards, S. J., Altig, R. A. & Gascon, C. (1994). Standard techniques for inventory and monitoring. In R. W. Heyer, M. A. Donnelley, R. W. McDiarmid, L. C. Hayek & M. S. Foster Eds.). *Measuring and monitoring biological diversity: standard methods for amphibians.* (pp. 384). Washington - Smithsonian Press.

Shannon, C. E. (1948). A mathematical theory of communication. *Bell System Technical Journal,* 27, 379-423, 623-656.

Smallbone, L. T., Luck, G. W. & Wassens, S. (2011). Anuran species in urban landscapes: Relationships with biophysical, built environment and socio-economic factors. *Landscape and Urban Planning,* 101, 43-51.

Smulders, M., Nelson, T. A., Jelinski, D. E., Nielsen, S. E. & Stenhouse, G. B. (2010). A spatially explicit method for evaluating accuracy of species distribution models. *Diversity and Distributions,* 16, 996-1008.

Soberón, J. & Peterson, A. T. (2005). Interpretation of models of fundamental ecological niches and species' distributional areas. *Biodiversity Informatics,* 2, 1-10.

Sokal, R. R. & Rohlf, R. F. (Eds.). (1995). *Biometry: The principles and practice of statistics in biological research.* New York - W.H. Freeman and Company.

Southerland, M. T., Jung, R. E., Baxter, D. P., Chellman, I. C., Mercurio, G. & Volstad, J. H. (2004). Stream salamanders as indicators of stream quality in Maryland, USA. *Applied Herpetology,* 2, 23-46.

Thullen, J. S., Sartoris, J. J. & Walton, W. E. (2002). Effects of vegetation management in constructed wetland treatment cells on water quality and mosquito production. *Ecological Engineering,* 18, 441-457.

Trotta-Moreu, N. & Lobo, J. M. (2010). Deriving the species richness distribution of Geotrupinae (Coleoptera: Scarabaeoidea) in Mexico from the overlap of individual model predictions. *Environmental Entomology,* 39, 42-49.

Tzoulas, K., Korpela, K., Venn, S., Yli-Pelkonen, V., Kazmierczak, A., Niemela, J. & James, P. (2007). Promoting ecosystem and human health in urban areas using green infrastructure: a literature review. *Landscape and Urban Planning,* 81, 167-178.

van den Berg, A. E., Hartig, T. & Staats, H. (2007). Preference for nature in urbanized societies: stress, restoration, and the pursuit of sustainability. *Journal of Social Issues,* 63, 79-96.

Vasconcelos, T., Rodriguez, M. A. & Hawkins, B. A. (2011). Species distribution modelling as a macroecological tool: a case study using New World amphibians. *Ecography,* 34, 1-10.

Waldron, J. L., C.K. Dodd, J. & Corser, J. D. (2003). Leaf litterbags: factors affecting capture of stream-dwelling salamanders. *Applied Herpetology,* 1, 23-36.

Welsh, H. H., Jr. & Droege, S. (2001). A case for using plethodontid salamanders for monitoring biodiversity and ecosystem integrity of North American forest. *Conservation Biology,* 15, 558-569.

Welsh, H. H., Jr. & Ollivier, L. M. (1998). Stream amphibians as indicators of ecosystem stress: a case study from California's redwoods. *Ecological Applications,* 8, 1118-1132.

Weyrauch, S. L. & Grubb Jr, T. C. (2004). Patch and landscape characteristics associated with the distribution of woodland amphibians in an agricultural fragmented landscape: an information-theoretic approach. *Biological Conservation,* 115, 443-450.

Whiles, M. R., Lips, K. R., Pringle, C. M., Kilham, S. S., Bixby, R. J., Brenes, R., Connelly, S., Colon-Gaud, J. C., Hunte-Brown, M., Huryn, A. D., Montgomery, C. & Peterson, S. (2006). The effects of amphibian population declines on the structure and function of neotropical stream ecosystems. *Frontiers in Ecology and the Environment,* 4, 27-34.

Wisz, M. S., Hijmans, R. J., Li, J., Peterson, A. T., Graham, C. H., Guisan, A. & Distribut, N. P. S. (2008). Effects of sample size on the performance of species distribution models. *Diversity and Distributions,* 14, 763-773.

Wu, J., Jenerett, G. D., Buyantuyev, A. & Redman, C. L. (2011). Quantifying spatiotemporal patterns of urbanization: the case of the two fastest growing metropolitan regions in the United States. *Ecological Complexity,* 8, 1-8.

Zhang, Q. G. & Zhang, D. Y. (2007). Consequences of individual species loss in biodiversity experiments: an essentiality index. *Acta Oecologica,* 32, 236-242.

Table 1. Estimates of how much SDMs over-predicted species richness values in green spaces in and around Hamilton County, Ohio. Values in the table represent the percentage each threshold (MTP = minimum training presence threshold, MTSPS = maximum training sensitivity plus specificity threshold, 10% TP = 10 percentile training presence threshold, FC10 = fixed cumulative value 10 threshold) over-predicted species richness compared to surveyed values for all sites (overall), urban sites (sites with ≥ 50% urbanization within the 2-km buffer of centroid), non-urban sites (sites with ≤ 50% urbanization within 2-km buffer of centroid), and from eight sites where past species richness values (derived from county-park surveys) were added to our current species richness values derived from current surveys (labeled Additional).

Table 2. Summary species richness data for amphibians at sites sampled ≥ 10 times in the Hamilton County, Ohio area.

Table 3. Results from a general linear model investigating the factors that influenced the observed amphibian species richness within each site and associated parameter estimates and 95% confidence intervals (CI). Percentage data were square root transformed and number of wetlands within 2-km buffer was log transformed. Excluded correlated variables not chosen for analysis were as follows: size; standard deviation of land cover within parks and 2-km buffer of centroid; percent forest, non-forest natural, and urban cells within 2-km buffer of centroid, and number of aquatic habitats surveyed within each site.

Table 4. Best Akaike Information Criteria (AIC) models for observed species richness across all 20 green spaces. Shown are only the confidence set of candidate models (i.e., models with AIC w_i within 10% of highest model). Variables included were percent urbanization within 2-km buffer, number of wetlands within 2-km buffer, percent forest within site, and percent non-forest natural habitat within each site.

Table 1. Estimates of how much SDMs over-predicted species richness values in green spaces in and around Hamilton County, Ohio. Values in the table represent the percentage each threshold (MTP = minimum training presence threshold, MTSPS = maximum training sensitivity plus specificity threshold, 10% TP = 10 percentile training presence threshold, FC10 = fixed cumulative value 10 threshold) over-predicted species richness compared to surveyed values for all sites (overall), urban sites (sites with ≥ 50% urbanization within the 2-km buffer of centroid), non-urban sites (sites with ≤ 50% urbanization within 2-km buffer of centroid), and from eight sites where past species richness values (derived from county-park surveys) were added to our current species richness values derived from current surveys (labeled Additional).

Urbanization Category	MTP	FC10	10% TP	MTSPS
Overall ($n = 20$)	558	467	373	253
Non-urban ($n = 6$)	619	572	468	343
Urban ($n = 14$)	532	422	333	215
Additional ($n = 8$)	98	81	50	14

Table 2. Summary species richness data for amphibians at sites sampled ≥ 10 times in the Hamilton County, Ohio area.

Site	S_{obs} [a]	Chao 2 est. richness [b]	95% lower CI [b]	95% upper CI [b]	% Coverage [c]	% Exclusive species [d]	% Completeness [e]
Miami	12	12	12	12	100	10	60
Shawnee	10	10	10	11	100	0	50
Farbach	5	10	6	46	53	0	25
Triple Creek	9	9	9	9.8	100	0	45
Richardson	14	14	14	20	98	5	70
Withrow	9	9	9	10	100	15	45

a Number of species observed.

b Number of species estimated to be present (and associated 95% confidence intervals) based on the Chao 2 estimator.

c Number of species observed as a percentage of the average estimated species richness (Chao 2).

d Number of species not found elsewhere as a percentage of the landscape total.

e Number of species observed as a percentage of the landscape total.

Table 3. Results from a general linear model investigating the factors that influenced the observed amphibian species richness within each site and associated parameter estimates and 95% confidence intervals (CI). Percentage data were square root transformed and number of wetlands within 2-km buffer was log transformed. Excluded correlated variables not chosen for analysis were as follows: Size; standard deviation of land cover within parks and 2-km buffer of centroid; percent forest, non-forest natural, and urban cells within 2-km buffer of centroid, and number of aquatic habitats surveyed within each site.

Source	df	MS	F	P	Parameter Estimate	-95% CI	+95% CI
% urbanization within 2-km buffer	1	11.717	5.465	0.034	-5.66	-10.82	-0.50
No. wetlands within 2-km buffer	1	15.118	7.052	0.018	0.97	0.19	1.75
% forest within site	1	0.593	0.277	0.607	0.79	-2.40	3.98
% non-forest natural within site	1	0.111	0.052	0.823	0.52	-4.35	5.39
Error	15	2.144	–	–	–	–	–

Table 4. Best Akaike Information Criteria (AIC) models for observed species richness across all 20 green spaces. Shown are only the confidence set of candidate models (i.e., models with AIC w_i within 10% of highest model). Variables included were percent urbanization within 2-km buffer, number of wetlands within 2-km buffer, percent forest within site, and percent non-forest natural habitat within site.

Model	AICc	Δ AICc	w_i
No. Wetlands + % Urbanization	72.19	0.00	0.61
No. Wetlands + % Urbanization + % Forest	75.35	3.16	0.13
No. Wetlands + % Urbanization + % Non-forest	75.35	3.16	0.13
No. Wetlands + % Non-forest	76.26	4.07	0.08

Figure 1. (a) Distribution of locality points (black dots) used to model species suitable habitat within the watershed boundary. The boundary is located across several states with Ohio and Kentucky (the states where the surveys were conducted) located in the northeast and southern border of the map; and (b) the three counties where each green space (site) was surveyed; metropolitan areas are outlined with a red line and surveyed green spaces are represented by filled green polygons. Surveys were conducted throughout various locations within each site. The four lower panels represent the predicted amphibian species richness maps within the three counties surveyed as derived from the maximum training sensitivity plus specificity threshold (MTSPS) (c), minimum training presence threshold (MTP) (d), 10 percentile training presence threshold (10% TP) (e), and fixed cumulative value 10 threshold (FC10) (f)

Figure 2. Sample-based species accumulation curves for the six heavily sampled sites in the Hamilton County, Ohio metropolitan area

Appendix A. Area under curve (AUC) values.

Appendix B. Species richness maps.

Appendix C. Summary data.

Appendix D. Results from analysis of covariance.

Appendix A. Area under curve (AUC) values for each species modeled.

Species	AUC	Sample size
Northern Cricket Frog (*Acris crepitans*)	0.776	89
Streamside Salamander (*Ambystoma barbouri*)	0.816	31
Jefferson Salamander (*Ambystoma jeffersonianum*)	0.808	48
Spotted Salamander (*Ambystoma maculatum*)	0.827	54
Marbled Salamander (*Ambystoma opacum*)	0.957	28
Small-mouthed Salamander (*Ambystoma texanum*)	0.932	20
American Toad (*Bufo americanus*)	0.738	121
Fowler"s Toad (*Bufo fowleri*)	0.782	23
Northern Dusky Salamander (*Desmognathus fuscus*)	0.831	20
Southern Two-lined Salamander (*Eurycea cirrigera*)	0.785	58
Long-tailed Salamander (*Eurycea longicauda*)	0.868	20
Cave Salamander (*Eurycea lucifuga*)	0.838	27
Spring Salamander (*Gyrinophilus porphyriticus*)	0.948	37
Cope"s Grey Tree Frog (*Hyla chrysoscelis*)	0.799	65
American Bullfrog (*Rana catesbeiana*)	0.701	45
Green Frog (*Rana clamitans*)	0.736	114
Northern Leopard Frog (*Rana pipiens*)	0.798	40
Wood Frog (*Rana sylvatica*)	0.887	23
Eastern Red-backed Salamander (*Plethodon cinereus*)	0.828	37
Northern Ravine Salamander (*Plethodon electromorphus*)	0.919	20

Northern Slimy Salamander (*Plethodon glutinosus*)	0.901	21
Spring Peeper (*Pseudacris crucifer*)	0.778	99
Western Chorus Frog (*Pseudacris triseriata*)	0.883	24

Appendix B. Depictions of: a) Cities of Indianapolis, Indiana (1), Cincinnati, Ohio (2) Lexington, Kentucky (3), and Huntington, West Virginia (4) located within the watershed boundary. Predicted amphibian species richness maps within the watershed boundary derived from the maximum training sensitivity plus specificity threshold (b), minimum training presence threshold (c), 10 percentile training presence threshold (d), and fixed cumulative value 10 threshold (e).

Appendix C. Summary data including size (ha), species richness (SR; observed [Obs.] and predicted), and landscape variables (within 2-km buffer of the centroid of each site and within each site) for each site. MTP = minimum training presence threshold, MTSPS = maximum training sensitivity plus specificity threshold, 10% TP = 10 percentile training presence threshold, and FC10 = fixed cumulative value 10 threshold. FRS = Forested area, NFRS = Non-forested natural area, URB = urban area.
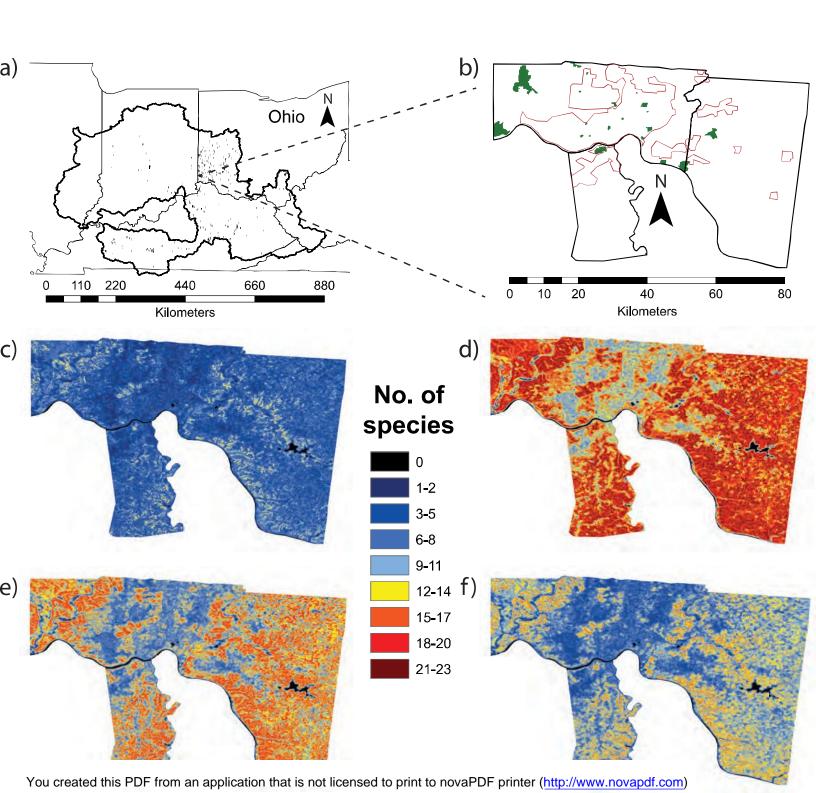
Site	Size (ha)	Obs. SR	FC10	MTP	MTSPS	10% TP	% in 2-km buffer FRS	URB	NFRS	% in site FRS	URB	NFRS	No. wetlands within buffer	No. aquatic habitats surveyed
					SR threshold						% in site			
Alms Park	38	1	17	21	13	17	26.7	64.5	6.6	81.4	16.2	2.4	6	0
Ault Park	89	2	20	23	14	20	23.4	57.9	7.6	76.4	21.4	2.3	6	2
Avon Woods	15	1	16	22	10	16	6.1	92.1	1.7	63.6	6.9	29.5	1	2
Bracken Woods	11	3	16	22	9	16	43.0	55.8	0.8	90.9	9.1	0.0	1	1
Burnet Woods	41	0	15	19	9	15	10.4	89.0	0.4	38.1	58.4	1.3	1	0
Cincinnati Nature Center	414	9	22	23	14	22	67.6	17.8	13.0	93.2	1.6	4.8	33	6
Devou Park	324	2	21	23	13	21	35.4	55.4	2.5	69.6	28.4	1.5	9	2
Farbach-Werner	9	5	14	17	7	14	16.0	83.6	0.3	72.7	27.3	0.0	24	3
French Park	115	2	20	23	13	20	25.6	72.8	1.4	83.3	15.7	0.8	6	2
Glenway Woods	8	2	18	21	10	18	26.2	73.1	0.1	98.7	1.3	0.0	2	0
Kennedy Park	3	1	11	16	7	11	19.3	79.4	1.4	34.4	65.6	0.0	2	1

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Site													
Koeing Park	5	0	12	14	7	12	7.7	91.6	0.2	0.0	100	0.0	4	0
Miami Forest	1758	9	22	23	15	22	53.4	7.9	35.8	54.4	6.9	36.0	13	4
Rapid Run Park	22	1	16	20	8	16	16.1	83.4	0.5	50.6	46.9	4.9	1	1
Richardson	159	5	21	23	14	21	60.1	21.5	17.1	71.0	7.7	21.3	24	4
Seymour	83	2	22	23	13	22	29.5	65.0	4.4	87.0	6.1	6.9	5	2
Shawnee Forest	613	6	21	23	14	21	40.7	4.2	45.2	45.2	1.8	43.5	11	5
Triple Creek Park	72	5	16	19	8	16	30.1	66.5	2.8	16.8	72.1	4.5	21	2
Withrow	109	3	21	23	14	21	45.6	32.9	5.2	94.1	2.2	3.7	4	2
Woodland Mounds	431	5	22	23	14	22	68.3	13.4	7.0	80.4	9.2	10.4	19	3

Appendix D. Results of ANCOVA and regression analyses to determine if species richness observations/estimates differed across the urban gradient. Species richness was the dependent variable, which was estimated either through field-observations (observed SR) or Maxent-predictions (predicted SR). Square root transformed percent urbanization within 2 km of site was the covariate in the analysis. The non-significant interaction term indicates that the relationship between species richness estimates and urbanization was not different. Estimates for each method differed significantly, and urbanization had a significant, negative effect on richness estimates. Linear equations for observed and predicted species richness and percent urbanization within 2-km are as follows: observed species richness, $r^2 = 0.627$, $P \leq 0.001$, Observed species richness = $9.597 – 0.896*\%$ urbanization; predicted species richness, $r^2 = 0.491$, $P \leq 0.001$, Predicted species richness = $23.765 – 0.925*\%$ urbanization.

Source	df	MS	F	P
Predicted SR	1	1949.510	521.920	≤ 0.0001
% urbanization within 2-km buffer	1	180.590	48.348	≤ 0.0001
Predicted SR * % urbanization within 2-km buffer	1	0.05	0.0121	0.913
% forest within site	36	3.740	–	–

Figure 1.

Appendix A.

Acknowledgments