

Article



Predation Pressure on Sentinel Insect Prey Along a Riverside Urbanization Gradient in Hungary

Csaba Béla Eötvös 1,2,*, Gábor L. Lövei ³ and Tibor Magura ¹

- ¹ Department of Ecology, University of Debrecen, H-4032 Debrecen, Hungary; maguratibor@gmail.com
- ² Department of Forest Protection, NARIC Forest Research Institute, H-3232 Mátrafüred, Hungary
- ³ Department of Agroecology, Aarhus University, Flakkebjerg Research Centre, DK-4200 Slagelse, Denmark; gabor.lovei@agro.au.dk
- * Correspondence: eotvos.csaba@erti.naik.hu; Tel.: +36-30-382-0375

Received: 8 January 2020; Accepted: 30 January 2020; Published: 1 February 2020

Abstract: Urbanization is one of the most important global trends which causes habitat reduction and alteration which are, in turn, the main reasons for the well-documented reduction in structural and functional diversity in urbanized environments. In contrast, effects on ecological mechanisms are less known. Predation is one of the most important ecological functions because of its community-structuring effects. We studied six forest habitats along a riverside urbanization gradient in Szeged, a major city in southern Hungary, crossed by the river Tisza, to describe how extreme events (e.g., floods) as primary selective pressure act on adaptation in riparian habitats. We found a generally decreasing predation pressure from rural to urban habitats as predicted by the increasing disturbance hypothesis (higher predator abundances in rural than in urban habitats). The only predators that reacted differently to urbanization were ground active arthropods, where results conformed to the prediction of the intermediate disturbance hypothesis (higher abundance in moderately disturbed suburban habitats). We did not find any evidence that communities exposed to extreme flood events were preadapted to the effects of urbanization. The probable reason is that changes accompanied by urbanization are much faster than natural landscape change, so the communities cannot adapt to them.

Keywords: urban; rural; ecological function; predation; predation paradox; sentinel prey; caterpillar; arthropod; mammal; bird

1. Introduction

Urbanization is as old as the first cities which appeared between 5100 BC and 2900 BC in the Fertile Crescent [1]. Today, urbanization is one of the most important processes shaping our environment, with fewer people living in rural than urban areas globally [2]. By 2050, the global rural population is expected to be ca. 3.1 billion, slightly less than today, while the urban populations are projected to reach 6.7 billion [2]. Globally, urbanization has several similar elements. From rural areas to urban centres, the original habitat matrix becomes smaller and more fragmented, road densities increase, along with the area covered by artificial surfaces, with air and soil pollution often showing the same trend [3,4]. Other changes include more human disturbance, increased noise level, and changes in temperature and precipitation patterns [5].

The biodiversity of the area affected by urbanization changes significantly. The urban matrix is rarely suitable for most of the original inhabitants of the rural habitat [6,7], so there is a reorganization of biodiversity [8,9]. Besides the changes in habitat amount, distribution and quality, new sources of mortality—including novel xenobiotics, hunting, collision with structures or vehicles, electrocution, and predation in the urban environment—can decrease survival rates [10]. Hence, several animal and

plant species present in the original rural habitat decrease in density or disappear altogether from urban habitats [11] and are often replaced by non-native species [12,13]. Effects of urbanization can also cause several physiological and behavioural changes in body size [14], body size distribution [15], fluctuating asymmetry [16], migratory behaviour [17], as well as lower reproduction [18] and survival rates [19]. Habitat specialists often disappear from urbanized areas [20–22] and consequently, functional diversity can decrease [23]. However, urbanization does not necessarily result in losses in taxonomic diversity or species richness [24]. More importantly, urbanization can change ecological functioning such as biogeochemical cycles [25] and trophic interactions (pollination [26], parasitism [27], predation [28,29]).

Predation, due to its community structuring effects, is one of the most important ecological processes [30]. Predator assemblages of urban habitats are different from those in rural ones [31,32]. Many predators avoid urban habitats, at least during daytime [33]. Populations of synanthropic predators can reach higher abundance in urban environments [31]. Local prey distribution can be changed by prey aggregating near light sources [34] or bird feeders [35]. Predator populations react to this [36], causing increased local predation rates [37]. Reaction to altered predation risk can lead to changes in behaviour [38], demographics, and interspecies interactions [39]. Current evidence indicates that predation pressure is lower in urban than rural areas, but this is mostly based on data on vertebrate predators; there is very little quantitative information about invertebrate predation [28].

Quantification of predation on invertebrate prey is difficult, because the attacks are mostly cryptic and evidence is difficult to obtain. Visual or video surveillance is complicated and expensive, and the activity of the predator may be affected by the observer or the equipment [40]. Gut content analysis or prey labelling produces results of varying resolution and secondary predation and the spread of the label can be related to non-predatory events [41]. Sentinel prey is one widely used method to measure predation intensity [42]. The prey can be immobile stages of arthropods such as eggs [43], pupae [44], or immobilized insects (e.g., aphids glued on self-adhesive paper [45]). The use of real prey is an advantage, but the identity of the predator usually remains unknown [42]. In contrast, artificial prey are not removed and the attack marks left by predators allow identification [46]. This method is suitable to compare predation pressure in various habitats [46].

We used the sentinel prey method to characterize predation on artificial caterpillars along an urbanization gradient in Szeged, Hungary. The specialty of this location is that this city—similarly to all the 50 largest built-up urban areas [47] except Mexico City—is built next to water. In spite of this feature, studies on ecological mechanisms in cities do not commonly consider this important factor. According to the natural flow-regime paradigm [48,49], extreme events (e.g., flood) exert a strong environmental filter in riparian habitats. With this in mind, we tested the following hypotheses:

• H1—The drastic, frequent floods constitute such a powerful environmental filter that other effects of urbanization are overwritten. Thus, there would be no difference in predation pressure along the gradient because floods equally affect all stages of the urbanization gradient.

• H2—Predation rates are lower in more urbanized habitats than in the rural ones. According to the increasing disturbance hypothesis [50], predator abundance decreases with advancing urbanization, and thus predation pressure would also decrease.

• H3—Bird predation will show a peak during breeding time. According to the match/mismatch hypothesis [51] the reproductive success of birds is maximized when they synchronize their reproduction with the peak of the food supply [52], around mid-May in Hungary [53].

• H4—Predation by small mammals will peak at the end of the growing season. The reproductive cycle of small mammals such as the wood mouse, *Apodemus sylvaticus* (L., 1758), in Europe shows its minimum in winter and its maximum in summer, producing the lowest population abundance in spring and summer, while the highest population abundances are in autumn and winter [54]. Consequently, we can expect the highest predation activity by small mammals during the autumn study period.

We found the highest predation rates in rural habitats by all predator groups except arthropods at ground level, refuting H1 while partially supporting H2: the effects of urbanization can overwrite

the effects of regular floods. Predation activity by birds and arthropods was the highest in spring and summer (H3 supported), while the highest peak of predatory activity by small mammals was recorded in summer (H4 not supported).

2. Materials and Methods

Our study site was in and around the city of Szeged (46°15′N; 20°8′E), 170 km southeast of Budapest (Hungary). This city lies on both sides of the lower reaches of the Tisza River. Upstream but within the city limits, it also receives the Maros River: both of them collect water from the Carpathians, and regularly flood their forested floodbeds (although no flood was registered during the period of study) (Table A1.). The floodbed was transformed during the end of the 19th century so, that the drainage of the water is optimal and the flood effect reducing infrastructure, built in 1973, is far enough upstream (150 km) from the city to assume equal flood intensity across the urban gradient [55]. The riverside forests are nearly continuous on the left bank, and more fragmented on the right one, allowing us to choose an urbanization gradient fulfilling the Globenet protocol conditions [56] (Figure 1).



Figure 1. The map of the study sites. A and B rural, C and D suburban, and E and F urban sites.

Vegetation originally was a primary forest of white willow (*Salix alba*) and white poplar (*Populus alba*). In the last decades, two invasive tree species [57], green ash (*Fraxinus pennsylvanica*) and box elder (*Acer negundo*) encroached on the area [55]. The undergrowth was dominated by invasive species, with a few natives (Table 1).

Common	Scientific name -	Number of dummy caterpillars on tree species in					Presence in undergrowth in						
name		R		S		U		R		S		U	
		1	2	1	2	1	2	1	2	1	2	1	2
Green ash*	Fraxinus pennsylvanica	24	1	30	16	7	4		Y		Y	Y	
Grey poplar	Populus x canescens	8	11	2	1	37	5				Y	Y	
White willow	Salix alba	15	3	8	6		32						
Box elder*	Acer negundo		25	7	7	2	4			Y	Y		Y
European White elm	Ulmus laevis	1	6	1	1	1							
Black poplar	Populus nigra		1		5	1							
White mulberry	Morus alba		1		3					Y			
Silver maple	Acer saccharinum				9		3				Y		
Field maple	Acer campestre										Y		
False indigo*	Amorpha fruticosa							Y	Y	Y	Y	Y	Y
Riverbank grape*	Vitis riparia									Y		Y	

Table 1. Tree and shrub species presence and the number of dummy caterpillars placed per session at the observed habitats on tree trunks. R-rural, S-suburban, U-urban, Y-given species is present at the location.

* Invasive in Hungary [57].

Data were collected during the growing seasons (April–October) in 2014–2016. Selected locations on both banks included rural, suburban, and urban areas, with an increase in the built-up area, increasing intensity of forest management, as well as visitation rates by city residents from rural to urban habitats (Figure 2). The average built up area (within a 500 m radius of a study site) was 0.3% in rural, 41.3% in suburban, and 54.8% in urban sites (Figure 2). In rural areas, cultivated fields dominated outside the dykes (12.5%) (Figure 2). In the riverside forest, the undergrowth was not managed in the rural sites, while in the urban sites the undergrowth was cut twice yearly. In the suburban areas, the last such operation was 2 years before the start of the study.

Along the two urbanization gradients, the average distance between sites was 5 km (range: 1–10 km). In each site, there were four patches with a minimum distance of 10 m between them. Within each patch, 12 trees were selected pseudo randomly by considering the species and trunk diameter. The average distance of selected trees within patches were 4.4 m (range: 0.5–23 m). On each of these, one dummy caterpillar was placed on the trunk and one at a random distance (range: 0–4 m) and direction from the base of the tree, on the ground. The dummy caterpillars (20 mm long, 3 mm thick) were made of light green plasticine (Smeedi plus, V. nr. 776609, Vilborg, Denmark), using a modified garlic press [46]. The colour, shape, and size imitated a general caterpillar prey [42]. The artificial prey was fixed to the bark of the trees or on a suitable surface on the ground with superglue (Pentack Super Glue, Pentacolor, Budapest, Hungary), and exposed for 24 h, then checked for attack marks using a handheld magnifying glass (10 ×). In case of doubt, the caterpillar was photographed and inspected on computer. Predators were identified by their characteristic marks left on the artificial prey (Figure 3). Attacks by different predators were considered independent events, but multiple marks by the same type of predator were classified as single attack. Overall, 12,672 caterpillars were exposed during the three years; 448 of them were not recovered.

For temporal analysis, May and June were considered spring, summer during July and August and autumn from September to the end of October.



Figure 2. Habitat types within a 500-meter radius of the study plots.



Figure 3. Characteristic marks left by different predator groups on dummy caterpillars. A–small mammals, B–arthropods, C–birds.

2.1. Data Analysis

We analysed data on ground-placed vs. trunk-placed caterpillars separately.

Before the start of the analysis, spatial autocorrelation was checked to decide whether the individual caterpillars could be considered independent, using Moran's I [58].

We used generalized linear mixed model (GLMM) [59,60] with urbanization stage and season as fixed effects, and study site as random factor. We found no significant differences among predation levels between the two gradients and the study years, so these factors were omitted.

Lognormal distribution fitted to our data best on the quantile to quantile plot, allowing us to use the penalized quasi-likelihood (PQL) method. PQL is a flexible technique that can deal with non-normal data, unbalanced design, and crossed random effects [61]. It effectively treats the random effects as 'fixed' and estimates them in a similar manner to other fixed effects as in a generalized linear model (GLM). Under a growing number of clusters, PQL estimates remain estimation consistent [62]. For multiple comparison of means we used Tukey test.

All calculation were made in R (version 3.5.1) [63]. Packages ncf [64] and lme4 [65] were used for autocorrelation methods. For GLMM calculations, we used car [66], MASS [67], and nlme [68].

Some of the marks were not related to predator attack, and in wet weather, snails occasionally left characteristic trails. In such case, the caterpillar was considered missing. A difference was considered significant at p < 0.05.

3. Results

There were 1780 (14.6%) attacks on sentinel prey. At ground level from the 6336 preys, 974 (16%) were attacked. On the soil surface small mammals were the most active predators (10.4%), followed by birds (3.4%), and arthropods (2.6%) (Table 2). From the trunk-placed 6336 caterpillars, 836 (13.2%) were attacked. On tree trunks, arthropods were the most active predators (8.8%), followed by small mammals (3.0%) and birds (1.2%) (Table 2). We were not able to identify predation marks on 448 (3.5%) dummy caterpillars because they were either missing or melted (Table 2).

Predation pressure (% prey attacked) on ground vs. trunk							
Group	Overall	Ground	Trunk				
-	(n = 12672)	(n = 6336)	(n = 6336)				
All predators	14.6	16.0	13.2				
Birds	2.3	3.4	1.2				
Mammals	6.7	10.4	3.0				
Arthropods	5.7	2.6	8.8				
Missing	3.5	5.4	1.6				

Table 2. Predation pressure by different predator groups

3.1. Predation Levels along the Urbanization Gradient

3.1.1. Overall Predation on Dummy Caterpillars

We found a significant, decreasing trend in attack frequency from rural to urban habitats both on trunk (rural - suburban: Estimate = -0.087, SD = 0.025, z = -3.555, p = 0.001, rural - urban: Estimate = 0.152, SD = 0.025, z = 6.098, p < 0.001, suburban - urban: Estimate = 0.065 SD = 0.025, z = 2.553, p = 0.029) and on ground level (rural - suburban: Estimate = -0.096, SD = 0.036, z = -2.700, p = 0.019, rural - urban: Estimate = 0.213, SD = 0.036, z = 5.921, p < 0.001, suburban - urban: Estimate = 0.118, SD = 0.036, z = 3.231, p = 0.004) (Figure 4).

3.1.2. Bird Predation on Dummy Caterpillars

There was a generally decreasing trend along the urbanization gradient (Figure 4). We found no significant differences in attack rates at ground level (rural - suburban: Estimate = -0.003, SD = 0.013, z = -0.241, p = 0.090, rural - urban: Estimate = 0.031, SD = 0.013, z = 2.342, p = 0.050, suburban - urban: Estimate = 0.028, SD = 0.013, z = 2.101, p = 0.969), but trunk-placed caterpillars in the rural habitat suffered significantly higher attack rates than those in the suburban or urban habitats (rural - suburban: Estimate = -0.014, SD = 0.006, z = -2.377, p = 0.046, rural - urban: Estimate = 0.022, SD = 0.006, z = 3.648, p < 0.001, suburban - urban: Estimate = 0.008, SD = 0.006, z = 1.271, p = 0.412) (Figure 4).

3.1.3. Mammalian Predation on Dummy Caterpillars

Similarly to birds, mammals showed a decreasing trend of predation activity along the urbanization gradient (Figure 4). However, mammal predation at ground level in rural habitats was significantly higher than in suburban or urban ones (rural - suburban: Estimate = -0.116, SD = 0.038, z = -3.067, p = 0.006, rural - urban: Estimate = 0.186, SD = 0.038, z = 4.906, p < 0.001, suburban - urban: Estimate = 0.071, SD = 0.038, z = 1.843, p = 0.156) (Figure 4). No significant difference was found on caterpillars placed on tree trunks (rural - suburban: Estimate = -0.035, SD = 0.028, z = -1.253, p = 0.422, rural - urban: Estimate = 0.059, SD = 0.028, z = 2.081, p = 0.094, suburban - urban: Estimate = 0.023, SD = 0.028, z = 0.828, p = 0.686) (Figure 4).

Arthropod attacks showed a decreasing trend along the urbanization gradient from rural to urban habitats on tree trunks but not on ground level (Figure 4). At ground level, suburban predation activity was significantly higher than in urban habitats, with a trend of higher activity in suburban than rural habitats (rural - suburban: Estimate = 0.019, SD = 0.009, z = 2.205, p = 0.070, rural - urban: Estimate = 0.007, SD = 0.009, z = 0.830, p = 0.685, suburban - urban: Estimate = 0.026, SD = 0.009, z = 3.035, p = 0.007) (Figure 4). On trunk placed prey, we found significantly decreasing predation levels from the rural to the urban habitats (rural - suburban: Estimate = -0.048, SD = 0.015, z = -3.260, p = 0.003, rural - urban: Estimate = 0.085, SD = 0.015, z = 5.628, p < 0.001, suburban - urban: Estimate = 0.037, SD = 0.015, z = 2.377, p = 0.046) (Figure 4).



Figure 4. Predation activity along the urbanization gradient on different placement and by different predator groups. Significance codes: ***: p < 0.001, **: p < 0.01, *: p < 0.05, $\therefore p < 0.1$.

3.2. Seasonal Trends

3.2.1. Overall Predation on Dummy Caterpillars

Ground level predators were most active during summer (spring - summer: Estimate = 0.087, SD = 0.017, z = 5.078, p < 0.001, spring - autumn: Estimate = -0.057, SD = 0.019, z = -3.105, p = 0.005, summer - autumn: Estimate = -0.144, SD = 0.018, z = -7.881, p < 0.001) while on tree trunks, there was significantly lower activity during autumn (spring - summer: Estimate = 0.018, SD = 0.016, z = 1.151, p = 0.482, spring - autumn: Estimate = -0.117, SD = 0.017, z = -6.918, p < 0.001, summer - autumn: Estimate = -0.135, SD = 0.017, z = -7.824, p < 0.001) (Figure 5).

3.2.2. Bird Predation on Dummy Caterpillars

Autumn predation levels were significantly lower than during spring or summer at both trunk (spring - summer: Estimate = -0.006, SD = 0.004, z = -1.620, p = 0.237, spring - autumn: Estimate = -0.020, SD = 0.004, z = -5.688, p < 0.001, summer - autumn: Estimate = -0.014, SD = 0.004, z = -3.942, p < 0.001) and ground level (spring - summer: Estimate = 0.006, SD = 0.008, z = 0.823, p = 0.689, spring - autumn: Estimate = -0.039, SD = 0.008, z = -5.007, p < 0.001, summer - autumn: Estimate = -0.045, SD = 0.008, z = -5.649, p < 0.001) (Figure 5).

3.2.3. Mammalian Predation on Dummy Caterpillars

Mammal predation activity was highest during the summer months both at ground level (spring - summer: Estimate = 0.087, SD = 0.014, z = 6.018, p < 0.001, spring - autumn: Estimate < 0.001, SD = 0.015, z = 0.025, p = 1, summer - autumn: Estimate = -0.086, SD = 0.015, z = -5.786, p < 0.001) and on the trunks (spring - summer: Estimate = 0.018, SD = 0.007, z = 2.669, p = 0.021, spring - autumn: Estimate < -0.001, SD = 0.007, z = -0.038, p = 0.999, summer - autumn: Estimate = -0.018, SD = 0.018, SD = 0.007, z = -2.619, p = 0.024) (Figure 5). After closer examination the peak predation was between August and September, but the autumn decrease was steeper, as in the case of birds.

3.2.4. Arthropod Predation on Dummy Caterpillars

Arthropod predators were less active during autumn than at other seasons, both at ground level (spring - summer: Estimate = 0.002, SD = 0.006, z = 0.293, p = 0.954, spring - autumn: Estimate = -0.025, SD = 0.006, z = -4.155, p < 0.001, summer - autumn: Estimate = -0.027, SD = 0.006, z = -4.310, p < 0.001) and on the trunks (spring - summer: Estimate = 0.005, SD = 0.014, z = 0.341, p = 0.938, spring - autumn: Estimate = -0.103, SD = 0.015, z = -6.745, p < 0.001, summer - autumn: Estimate = -0.108, SD = 0.016, z = -6.880, p < 0.001) (Figure 5).



Figure 5. Predation activity during the season on different placement and by different predator groups. Significance codes: ***: p < 0.001, **: p < 0.01, *: p < 0.05, :: p < 0.1.

4. Discussion

We found a generally decreasing predation activity along our urbanization gradient from rural to urban habitats except for ground-active arthropod predators (H2). This conforms to other findings [28,29] and does not indicate that communities regularly exposed to extreme flood events were insensitive to urbanization (H1). Urbanization can be a fast process, proceeding at a higher speed than natural landscape and environmental changes, creating highly dynamic and complex habitats [69]. The increasing disturbance hypothesis was supported by our data, except for arthropods at ground level, where Connell's intermediate disturbance hypothesis [70] seemed a more acceptable explanation, predicting the increase in predation pressure at intermediate levels of disturbance, in the studied situation at the suburban sites. Alternatively, the "predation paradox" hypothesis [28,71] can also explain the decreasing predation pressure toward cities, but we have no information on the abundances of the different predator groups.

Ground beetles constitute one of the main arthropod predator group on the soil surface, together with spiders and ants [72,73]. We found traces of attack by ground beetles in this experiment. The majority of previous studies considering ground beetle abundance do not support the intermediate disturbance hypothesis [13,74,75]. Thus, there can be differences in movement activity (reflected by pitfall trap catches) and foraging activity (attacks on sentinel prey) in this group. Nevertheless, in the same study area, spiders also show a similar pattern which can be explained by Connell's intermediate disturbance hypothesis [76] and ground beetle abundance was highest in rural habitats, moderate in suburban areas, and lowest in urban ones (S. Mizser, University of Debrecen, Debrecen, Hungary. Personal Communication, 2020.).

We found higher predation rates by birds in spring and summer than autumn. This result conformed to the match/mismatch hypothesis [51] (H3), except for the high predation rate during summer. This can be the result of the higher population densities just after the fledging of the nestlings. Slight differences in the date of laying the first egg (one week earlier in urban than rural habitats) and of the occurrence of peak caterpillar biomass (4 days later in urban habitats) [77] can slightly shift the results. For mammals, late summer was the most active period (H4). So, the population growth was not enough to increase the predation rate, but the increased protein needs during the reproduction period [78] can alter the eating habits of small mammals [79].

We found higher predation rates (14.6% overall, and 16.0% at ground level), than the previously reported 8.8% median predation rate on artificial prey [42]. This median was calculated from worldwide data and we have to consider that from the equator to the poles the predation activity in forested habitats gradually, but not significantly increases [42], so our result is not an outlier. Overall, we found higher predation rates at ground level which is also in agreement with previous results [42]. Vertebrate predation on artificial caterpillars (birds and mammals, 9.0% combined) was higher than arthropod predation (5.7%) as expected [42]. The same pattern was found at ground level (vertebrate predation 13.8%, arthropod predation 2.6%) but the opposite on tree trunk (vertebrate predation 4.2%, arthropod predation 8.8%). The higher arthropod predation above ground level is different from previously reported results [42]. This may reflect that higher small mammal activity at ground level probably included species that also prey on carabids and other ground-active arthropods, and their densities often show a negative relationship [80].

The urban predator assemblage is different from the natural and semi-natural communities [31]. Predator abundances and feeding habits are changing. For example, availability of anthropogenic food could, in the case of opportunistic predators, result in lower predation rates on certain taxa [81]. Other behavioural change can be triggered by more aggregated prey in urban habitats, for example around bird feeders [35] and light sources [34], which can result locally higher predation rates, and consequently lower predation activity at other places [37].

A rarely mentioned complication was that some caterpillars melted and had to be discarded – the soft plasticine cannot keep eventual attack marks. Initially, we attributed this to high

temperatures—during sunny summer days, the temperatures at the study site can reach >35 °C. However, this effect may have been due to exposure to direct sunshine rather than high temperatures, because the same plasticine was successfully used in the Negev Desert in Israel, also at temperatures well above 40 °C (M. Ferrante, University of the Azores, Terceira, Portugal. Personal Communication, 2020).

In this riverside forested landscape, we found the same pattern, a decreasing predation pressure from rural to urban habitats, as earlier studies in non-riverside urbanized areas on birds as prey [28], but it is not clear that the changes in the riparian community composition were the same as in the other forest communities in other cities. Thus, it can be useful to compare riparian and non-riparian habitats in the same city in order to possibly separate these confounding effects. It would be profitable to investigate different prey groups, not only insects, but also amphibians, reptiles, mammals, and birds at the same time and place.

5. Conclusions

In conclusion, we found a decreasing predation pressure on artificial insect prey along the urbanisation gradient from rural to urban forest, with different predators being the most active at ground level vs. tree trunk. Our general attack rates tended to be higher than the median values in the literature. Our studied gradient was special in that it was in a floodbed crossing a large city. Apparently, the floods did not constitute a strong enough habitat filtering effect to overwrite the impact of habitat changes brought by urbanisation.

Author Contributions: Concept and study design: G.L., C.E., T.M.; Data collection C.E.; Analysis C.E., T.M., G.L.; Writing shared by all authors. Author sequence is the "sequence by credit" principle.

Acknowledgments: This paper is in partial fulfilment of the PhD requirements at the University of Debrecen (CE). This publication was supported by the National Research, Development, and Innovation Fund (OTKA K-128008 and OTKA K-131459). We thank Marco Ferrante and Szabolcs Mizser for information.

Conflicts of Interest: The authors declare no conflict of interest. "The funders had no role in the design of the study; in the collection, analyses, or interpretation of data; in the writing of the manuscript, or in the decision to publish the results".

Appendix A

Period	Floodbed inundated (occasions)	Average length of flood (days)	Inundation ≥50 cm (occasions)	Average length of ≥50 cm inundation (days)	Average max depth of inundation (cm)
Whole year	25	44.6 (range 1–95)	20	41.0 (range 9–78)	173.1 (range 5–449)
April– October	24	35.6 (range 4–77)	20	31.9 (range 6–73)	172.1 (range 23–449)

Table A1. Flooding history of river Tisza at Szeged between 1977–2016

References

- 1. Willoughby, P.R. The evolution of the ancient city: urban theory and the archaeology of the Fertile Crescent. *Can. Stud. Popul.* **2013**, *40*, 107–108.
- 2. United Nations. World Urbanization Prospects: The 2018 Revision; United Nations: New York, NY, USA, 2018.
- 3. Medley, K.E.; McDonnell, M.J.; Pickett, S.T.A. Forest-landscape structure along an urban-to-rural gradient. *Prof. Geogr.* **1995**, *47*, 159–168.
- 4. Pickett, S.T.A.; Cadenasso, M.L.; Grove, J.M.; Nilon, C.H.; Pouyat, R. V.; Zipperer, W.C.; Costanza, R. Urban ecological systems: linking terrestrial ecological, physical, and socioeconomic components of metropolitan areas. *Annu. Rev. Ecol. Syst.* **2001**, *32*, 127–157.
- Grimmond, C.S.B. Climate of cities. In *The Routledge Handbook of Urban Ecology*; Douglas, I., Goode, D., Houck, M., Wang, R., Eds.; Routledge: London, UK, 2015; pp. 103–119.

- 7. Niemelä, J. Ecology and urban planning. *Biodivers. Conserv.* 1999, *8*, 119–131.
- 8. Stracey, C.M.; Robinson, S.K. Does nest predation shape urban bird communities? In *Urban Bird Ecology and Conservation;* University of California Press: Berkeley, CA, USA, **2012**; pp. 49–70.
- 9. Magura, T.; Lövei, G.L.; Tóthmérész, B. Conversion from environmental filtering to randomness as assembly rule of ground beetle assemblages along an urbanization gradient. *Sci. Rep.* **2018**, *8*, 16992.
- 10. Newton, I. Population Limitation in Birds; Academic Press: New York, NY, USA, 1998; ISBN 9780080879239.
- 11. Marzluff, J.M.; Bowman, R.; Donnelly, R. A historical perspective on urban bird research: trends, terms, and approaches. In *Avian Ecology and Conservation in an Urbanizing World*; Marzluff, J.M., Bowman, R., Donnelly, R., Eds.; Springer US: Boston, MA, USA, **2001**; pp. 1–17.
- 12. McIntyre, N.E. Ecology of urban arthropods: a review and a call to action. *Ann. Entomol. Soc. Am.* **2000**, *93*, 825–835.
- 13. Tóthmérész, B.; Máthé, I.; Balázs, E.; Magura, T. Responses of carabid beetles to urbanization in Transylvania (Romania). *Landsc. Urban Plan.* **2011**, *101*, 330–337.
- 14. Weller, B.; Ganzhorn, J.U. Carabid beetle community composition, body size, and fluctuating asymmetry along an urban-rural gradient. *Basic Appl. Ecol.* **2004**, *5*, 193–201.
- 15. Magura, T.; Tóthmérész, B.; Lövei, G.L. Body size inequality of carabids along an urbanisation gradient. *Basic Appl. Ecol.* **2006**, *7*, 472–482.
- 16. Elek, Z.; Lövei, G.L.; Bátki, M. No increase in fluctuating asymmetry in ground beetles (Carabidae) as urbanisation progresses. *Community Ecol.* **2014**, *15*, 131–138.
- 17. Partecke, J.; Gwinner, E. Inceased sedentariness in European blackbirds following urbanization: a consequence of local adaptation? *Ecology* **2007**, *88*, 882–890.
- 18. Seress, G.; Bókony, V.; Pipoly, I.; Szép, T.; Nagy, K.; Liker, A. Urbanization, nestling growth and reproductive success in a moderately declining house sparrow population. *J. Avian Biol.* **2012**, *43*, 403–414.
- Riley, S.P.D.; Sauvajot, R.M.; Fuller, T.K.; York, E.C.; Kamradt, D.A.; Bromley, C.; Wayne, R.K. Effects of urbanization and habitat fragmentation on bobcats and coyotes in southern California. *Conserv. Biol.* 2003, 17, 566–576.
- 20. Devictor, V.; Julliard, R.; Couvet, D.; Lee, A.; Jiguet, F. Functional homogenization effect of urbanization on bird communities. *Conserv. Biol.* 2007, *21*, 741–751.
- 21. Liker, A.; Papp, Z.; Bókony, V.; Lendvai, Á.Z. Lean birds in the city: Body size and condition of house sparrows along the urbanization gradient. *J. Anim. Ecol.* **2008**, *77*, 789–795.
- 22. Lövei, G.L.; Magura, T.; Tóthmérész, B.; Ködöböcz, V. The influence of matrix and edges on species richness patterns of ground beetles (Coleoptera: Carabidae) in habitat islands. *Glob. Ecol. Biogeogr.* 2006, *15*, 283–289.
- Sacco, A.G.; Rui, A.M.; Bergmann, F.B.; Müller, S.C.; Hartz, S.M.; Sacco, A.G.; Rui, A.M.; Bergmann, F.B.; Müller, S.C.; Hartz, S.M. Reduction in taxonomic and functional bird diversity in an urban area in Southern Brazil. *Iheringia. Série Zool.* 2015, 105, 276–287.
- 24. Magura, T.; Lövei, G.L.; Tóthmérész, B. Does urbanization decrease diversity in ground beetle (Carabidae) assemblages? *Glob. Ecol. Biogeogr.* 2010, *19*, 16–26.
- 25. Grimm, N.B.; Hale, R.L.; Cook, E.M.; Iwaniec, D.M. Urban biogeochemical flux analysis. In *The Routledge Handbook of Urban Ecology*; Douglas, I., Goode, D., Houck, M., Wang, R., Eds.; Routledge: London, UK, **2015**; pp. 503–520.
- 26. Harrison, T.; Winfree, R. Urban drivers of plant-pollinator interactions. Funct. Ecol. 2015, 29, 879-888.
- Fenoglio, M.S.; Videla, M.; Salvo, A.; Valladares, G. Beneficial insects in urban environments: Parasitism rates increase in large and less isolated plant patches via enhanced parasitoid species richness. *Biol. Conserv.* 2013, *164*, 82–89.
- 28. Eötvös, C.B.; Magura, T.; Lövei, G.L. A meta-analysis indicates reduced predation pressure with increasing urbanization. *Landsc. Urban Plan.* **2018**, *180*, 54–59.
- 29. Ferrante, M.; Lo Cacciato, A.; Lövei, G.L. Quantifying predation pressure along an urbanisation gradient in Denmark using artificial caterpillars. *Eur. J. Entomol.* **2014**, *111*, 649–654.
- 30. Shochat, E.; Warren, P.S.; Faeth, S.H.; McIntyre, N.E.; Hope, D. From patterns to emerging processes in mechanistic urban ecology. *Trends Ecol. Evol.* **2006**, *21*, 186–191.

- 31. Haskell, D.G.; Knupp, A.M.; Schneider, M.C. Nest predator abundance and urbanization. In *Avian Ecology and Conservation in an Urbanizing World*; Marzluff, J.M., R., B., R, D., Eds.; Springer US: Boston, MA, USA, **2001**; pp. 243–258.
- 32. Sorace, A. High density of bird and pest species in urban habitats and the role of predator abundance. *Ornis Fenn.* **2002**, *79*, 60–71.
- 33. Tigas, L.A.; Van Vuren, D.H.; Sauvajot, R.M. Behavioral responses of bobcats and coyotes to habitat fragmentation and corridors in an urban environment. *Biol. Conserv.* **2002**, *108*, 299–306.
- 34. Gaston, K.J.; Bennie, J.; Davies, T.W.; Hopkins, J. The ecological impacts of nighttime light pollution: a mechanistic appraisal. *Biol. Rev.* **2013**, *88*, 912–927.
- Clergeau, P.; Savard, J.-P.L.; Mennechez, G.; Falardeau, G. Bird Abundance and Diversity along an Urban-Rural Gradient: A Comparative Study between Two Cities on Different Continents. *Condor* 1998, 100, 413– 425.
- 36. Scanlon, A.T.; Petit, S. Biomass and biodiversity of nocturnal aerial insects in an Adelaide City park and implications for bats (Microchiroptera). *Urban Ecosyst.* **2008**, *11*, 91–106.
- 37. Gering, J.C.; Blair, R.B. Predation on artificial bird nests along an urban gradient: predatory risk or relaxation in urban environments? *Ecography* **1999**, *22*, 532–541.
- Laundre, J.W.; Hernandez, L.; Ripple, W.J. The Landscape of Fear: Ecological Implications of Being Afraid. Open Ecol. J. 2010, 3, 1–7.
- 39. Avilés, J.M.; Bednekoff, P.A. How do vigilance and feeding by common cranes *Grus grus* depend on age, habitat, and flock size? *J. Avian Biol.* **2007**, *38*, 690–697.
- 40. Kistner, E.J.; Lewis, M.; Carpenter, E.; Melhem, N.; Hoddle, C.; Strode, V.; Oliva, J.; Castillo, M.; Hoddle, M.S. Digital video surveillance of natural enemy activity on *Diaphorina citri* (Hemiptera: Liviidae) colonies infesting citrus in the southern California urban landscape. *Biol. Control* **2017**, 115, 141–151.
- 41. Kidd, N.A.C.; Jervis, M.A. Population Dynamics. In *Insects as Natural Enemies*; Springe: Dordrecht, The Netherlands, **2007**; pp. 435–523.
- 42. Lövei, G.L.; Ferrante, M. A review of the sentinel prey method as a way of quantifying invertebrate predation under field conditions. *Insect Sci.* **2017**, *24*, 528–542.
- 43. Wilson, A.P.; Hough-Goldstein, J.A.; Vangessel, M.J.; Pesek, J.D. Effects of Varying Weed Communities in Corn on European Corn Borer, *Ostrinia nubilalis* (Hübner) (Lepidoptera: Crambidae), Oviposition, and Egg Mass Predation. *Environ. Entomol.* **2004**, *33*, 320–327.
- 44. Rowe, R.L.; Goulson, D.; Doncaster, C.P.; Clarke, D.J.; Taylor, G.; Hanley, M.E. Evaluating ecosystem processes in willow short rotation coppice bioenergy plantations. *GCB Bioenergy* **2013**, *5*, 257–266.
- 45. Östman, Ö. The relative effects of natural enemy abundance and alternative prey abundance on aphid predation rates. *Biol. Control* **2004**, *30*, 281–287.
- 46. Howe, A.; Lövei, G.L.; Nachman, G. Dummy caterpillars as a simple method to assess predation rates on invertebrates in a tropical agroecosystem. *Entomol. Exp. Appl.* **2009**, *131*, 325–329.
- 47. *Demographia World Urban Areas 14th Annual Edition;* Available online: http://www.demographia.com/db-worldua.pdf. (Accessed on 20 December 2019).
- 48. LeRoy Poff, N.; David Allan, J.; Bain, M.B.; Karr, J.R.; Prestegaard, K.L.; Richter, B.D.; Sparks, R.E.; Stromberg, J.C. The Natural Flow Regime A paradigm for river conservation and restoration. *Bioscience* **1997**, 47, 769–784.
- 49. Richter, B.D.; Baumgartner, J. V.; Powell, J.; Braun, D.P. A Method for Assessing Hydrologic Alteration within Ecosystems. *Conserv. Biol.* **1996**, *10*, 1163–1174.
- 50. Gray, J.S. Effects of environmental stress on species rich assemblages. Biol. J. Linn. Soc. 1989, 37, 19–32.
- 51. Cushing, D.H. Plankton production and year-class strength in fish populations: An update of the match/mismatch hypothesis. *Adv. Mar. Biol.* **1990**, *26*, 249–293.
- 52. Durant, J.; Hjermann, D.; Ottersen, G.; Stenseth, N. Climate and the match or mismatch between predator requirements and resource availability. *Clim. Res.* **2007**, *33*, 271–283.
- 53. Bereczki, K.; Ódor, P.; Csóka, G.; Mag, Z.; Báldi, A. Effects of forest heterogeneity on the efficiency of caterpillar control service provided by birds in temperate oak forests. *For. Ecol. Manage.* **2014**, *327*, 96–105.
- 54. Montgomery, W.I. Population structure and dynamics of sympatric Apodemus species (Rodentia: Muridae). *J. Zool.* **1980**, *192*, 351–377.

- 56. Elek, Z. Ground beetle (Coleoptera, Carabidae) assemblages along an urbanisation gradient near Sorø, Zealand, Denmark. *Entomol. Meddelelser* **2005**, *73*, 115–121.
- 57. Balogh, L.; Dancza, I.; Király, G. A magyarországi neofitonok időszerű jegyzéke, és besorolásuk inváziós szempontból. In *Biológiai inváziók Magyarországon Özönnövények*; TermészetBÚVÁR Alapítvány Kiadó: Budapest, Hungary, **2004**; pp. 61–92 ISBN 963 86107 5 1. (In Hungarian)
- 58. Moran, P.A.P. Notes on Continuous Stochastic Phenomena. *Biometrika* 1950, 37, 17–23.
- 59. Bates, D.M. lme4: Mixed-effects modeling with R. Springer, Berlin, Germany, 2010.
- Bolker, B.M.; Brooks, M.E.; Clark, C.J.; Geange, S.W.; Poulsen, J.R.; Stevens, M.H.H.; White, J.-S.S.; Henry, M.; Stevens, H.; White, J.-S.S. Generalized linear mixed models: a practical guide for ecology and evolution. *Trends Ecol. Evol.* 2009, 24, 127–135.
- 61. Breslow, N.E.; Clayton, D.G. Approximate Inference in Generalized Linear Mixed Models. J. Am. Stat. Assoc. 1993, 88, 9.
- 62. Vonesh, E.F.; Wang, H.; Nie, L.; Majumdar, D. Conditional Second-Order Generalized Estimating Equations for Generalized Linear and Nonlinear Mixed-Effects Models. *J. Am. Stat. Assoc.* **2002**, *97*, 271–283.
- 63. R Core Team R: A Language and Environment for Statistical Computing. Available online: https://www.R-project.org/ (Accessed on 29 December 2019).
- 64. Bjornstad, O.N.; Cai, J. Spatial Covariance Functions [R package ncf version 1.2-8] Available online: https://cran.r-project.org/package=ncf (accessed on 19/05/2019).
- Bates, D.; Mächler, M.; Bolker, B.; Walker, S. Fitting Linear Mixed-Effects Models Using Ime4. J. Stat. Softw. 2015, 67, 1–48.
- 66. Fox, J.; Weisberg, S. *An R Companion to Applied Regression*; Sage Publications: Thousand Oaks, CA, USA, **2018**; ISBN 9781544336473.
- 67. Venables, W.N.; Ripley, B.D. *Modern Applied Statistics with S*; Fourth Edition; Springer: New York, NY, USA, **2002**; ISBN 0387954570.
- 68. Pinheiro, J.; Bates, D.; DebRoy, S.; Sarkar, D.; R Core Team nlme: Linear and Nonlinear Mixed Effects Models. R package version 3.1-140 Available online: https://cran.r-project.org/package=nlme (accessed on 19/05/2019).
- 69. Antrop, M. Landscape change and the urbanization process in Europe. Landsc. Urban Plan. 2004, 67, 9–26.
- 70. Connell, J.H. Diversity in tropical rain forests and coral reefs. Science 1978, 199, 1302–1310.
- 71. Fischer, J.D.; Cleeton, S.H.; Lyons, T.P.; Miller, J.R. Urbanization and the Predation Paradox: The Role of Trophic Dynamics in Structuring Vertebrate Communities. *Bioscience* **2012**, *62*, 809–818.
- 72. Kajak, A.; Breymeyer, A.; Petal, J. Productivity investigation of two types of meadows in the Vistula Valley. XI. Predatory arthropods. *Ekol Pol Ser A* **1971**, 19, 223–233.
- 73. Nyffeler, M.; Birkhofer, K. An estimated 400–800 million tons of prey are annually killed by the global spider community. *Sci. Nat.* **2017**, *104*, 30.
- 74. Elek, Z.; Lövei, G.L. Patterns in ground beetle (Coleoptera: Carabidae) assemblages along an urbanisation gradient in Denmark. *Acta Oecologica* **2007**, *32*, 104–111.
- 75. Magura, T.; Tóthmérész, B.; Molnár, T. Changes in carabid beetle assemblages along an urbanisation gradient in the city of Debrecen, Hungary. *Landsc. Ecol.* **2004**, *19*, 747–759.
- 76. Tajthi, B.; Horváth, R.; Mizser, S.; Nagy, D.D.; Tóthmérész, B. Spider assemblages in floodplain forests along an urbanization gradient. *Community Ecol.* **2017**, *18*, 311–318.
- 77. Seress, G.; Hammer, T.; Bókony, V.; Vincze, E.; Preiszner, B.; Pipoly, I.; Sinkovics, C.; Evans, K.L.; Liker, A. Impact of urbanization on abundance and phenology of caterpillars and consequences for breeding in an insectivorous bird. *Ecol. Appl.* 2018, *28*, 1143–1156.
- Speakman, J.R. The physiological costs of reproduction in small mammals. *Philos. Trans. R. Soc. B Biol. Sci.* 2008, 363, 375–398.
- 79. Fitch, H.S. Seasonal Acceptance of Bait by Small Mammals. J. Mammal. 1954, 35, 39.
- 80. Parmenter, R.R.; MacMahon, J.A. Factors Influencing Species Composition and Population Sizes in a Ground Beetle Community (Carabidae): Predation by Rodents. *Oikos* **1988**, *52*, 350.

81. Contesse, P.; Hegglin, D.; Gloor, S.; Bontadina, F.; Deplazes, P. The diet of urban foxes (*Vulpes vulpes*) and the availability of anthropogenic food in the city of Zurich, Switzerland. *Mamm. Biol.* - *Zeitschrift für Säugetierkd.* **2004**, *69*, 81–95.



© 2020 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution

(CC BY) license (http://creativecommons.org/licenses/by/4.0/).