

1 On the lookout for danger: House Sparrow alert distance in three cities

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12
13 **Abstract** The House Sparrow *Passer domesticus*, closely associated with human presence along
14 urban–agricultural landscapes and widely distributed species outside its native range, has shown
15 great morphological and physiological variations, with its plasticity linked to its invasiveness.
16 Yet, there is a dearth of knowledge on the escape behavior of this sparrow along its North
17 American invasion range. We here assessed House Sparrows alert distances in Los Angeles and
18 Mexico City, but also in Barcelona as a control within its native distribution, considering ‘city’,
19 ‘sex/age’ (adult males vs. adult females and immatures), and ‘flock size’ as alert distance
20 predictors. Through a linear model and a classification and regression tree, we identified that
21 House Sparrow alert distances in Barcelona were significantly larger when compared to both
22 studied North American cities (Los Angeles, Mexico City). Given that alert distances were also
23 significantly larger in Los Angeles when contrasted with those from Mexico City, where it has

24 been recently shown to be hyper-abundant, our results also suggest that its boldness could be
25 related not only to an origin (native/non-native) driver, but also to a density-dependent process.

26

27 **Keywords** Barcelona, flushing distance, Los Angeles, Mexico City, *Passer domesticus*, urban
28 ecology.

29 **Introduction**

30 The establishment or growth of an urban system goes beyond the often simplistic view of the
31 replacement of pre-existing land-uses. Instead urbanization processes represent severe physical
32 and ecological changes associated with fulfilling modern urban needs (Eldredge and Horenstein
33 2014). Although urbanization filters regional avian species pools, limiting the presence of an
34 important proportion of species in cities (Aronson et al. 2014), there is growing evidence of
35 changes to wildlife associated with adaptive responses to urbanization by those species able to
36 overcome urban hazards and able to use the novel array of resources, which can ultimately drive
37 adaptation and evolution (Díaz et al. 2013; McDonnell and Hahs 2015; Johnson and Munshi-
38 South 2017; Samia et al. 2017). In the light of an increasingly urbanized planet, there is a
39 pressing need to understand wildlife responses to urbanization (McDonnell and Hahs 2015;
40 McDonnell and MacGregor-Fors 2016).

41 Fischer et al. (2015) suggested three types of urban wildlife species categories based on
42 their population response to urbanization: (1) ‘avoiders,’ (2) ‘utilizers,’ and (3) ‘dwellers.’
43 Among urban ‘dwellers,’ some species can experience population explosions, which Blair (1996)
44 coined as ‘exploiters.’ Avian urban exploiters have been widely studied in the search of the traits
45 related to the successful species facing such a dramatic environmental scenario as the urban one
46 (e.g., Kark et al. 2006; Vincze et al. 2015). Exploiters have also received attention in regards to
47 their potential negative effects with native urban avifauna (MacGregor-Fors et al. 2010,
48 González-Oreja et al. 2018). Among the most frequent cosmopolitan bird species, the Rock Dove
49 *Columba livia* and House Sparrow *Passer domesticus* head the list, with both of them considered
50 urban exploiters in many regions of the globe (Aronson et al. 2014).

51 The House Sparrow, which has been a human commensal species known to live in
52 Eurasia before modern times (Lowther and Cink 2006), has been subject to study for decades due
53 to its impressive range-expanding invasive capacity in several regions worldwide (Aronson et al.
54 2014). Currently, the House Sparrow has populations in most of Europe, a large proportion of
55 Western, South-Central, and South-Eastern Asia, South-East Oceania, Southern Africa, most
56 North America (except North-Central Canada and Greenland), Central America, and South
57 America (except the Amazon Basin and the Caribbean region) (Summers-Smith et al. 2018).
58 Specifically for North America, the House Sparrow was introduced in the New England area in
59 at least 16 independent events from 1850–1881 (Brown and Wilson 1975). After colonizing most
60 of the United States, it is presumed that the House Sparrow arrived to Mexico City by the 1930s
61 (Wagner 1959), only 49–80 years after its initial introduction. Studies focused on House Sparrow
62 morphological differences have found great variation across its North American range, generally
63 regarding its adaptation to contrastingly different environmental and climatic scenarios in the
64 face of the contrasting environmental heterogeneity of the regions it has successfully invaded
65 (Johnston and Selander 1964, 1971; Lowther 1977).

66 House Sparrows have been extensively studied along their North American invasion
67 range (mostly in Northern North America) for many decades now, including evidence of
68 morphological differences along their northern invasion range, as well as their physiology
69 (Johnston and Selander 1964, 1973; Kendeigh 1976, Lowther 1977, Martin et al. 2004).
70 Yet, there is an important dearth of knowledge on the behavior of this species, essentially
71 circumscribed to urban–agricultural landscapes throughout its North American invasion
72 distribution. Recently, a study focused on House Sparrow densities in three urban–agricultural
73 landscapes related to three cities (i.e., Barcelona, Los Angeles, Mexico City) showed differing

74 results among and within landscapes (MacGregor-Fors et al. 2017). Briefly, this study reports
75 higher densities in the studied landscape including Mexico City and lower densities in those of
76 Los Angeles and Barcelona, showing that their densities are not dependent from the origin of
77 their populations (i.e., native, exotic).

78 In this study, we assessed House Sparrow escape distances in the same three cities
79 considered in the aforementioned study (i.e., Barcelona, Los Angeles, Mexico City). We focused
80 on escape distances as they have shown to be a robust way to assess predation risk, specifically
81 the reaction to an approaching potential predator (Ydenberg and Dill 1986). In particular, alert
82 distance (i.e., distance at which birds become aware of a threat) has been used as an indicator of
83 the ability of birds to detect potential predators (Fernández-Juricic et al. 2001, Blumstein et al.
84 2004). Thus, escape distances have been long considered as useful approximations of how the
85 boldness or shyness of individuals (Wilson et al. 1994).

86 We had two mutually exclusive predictions related to a behavioral response (i.e., alert
87 distances) of House Sparrow and their population densities: (1) if their invasiveness is related
88 with their boldness, we then predicted individuals from Los Angeles and Mexico City (which are
89 part of the North American invasion population) to be less wary about human approaches than
90 those from Barcelona (which is part of the distribution range considered to be native for the
91 species), resulting in shorter alert distances in these two cities, and (2) if their densities are
92 related to their boldness, we then expected individuals in the city where higher densities were
93 recorded (i.e., Mexico City: average density during the breeding season: 11.6–21.7 ind/ha;
94 MacGregor-Fors et al. 2017) to be less wary about human approaches than those from lower
95 density cities (i.e., Barcelona: average density during the breeding season: 4.5–5.7 ind/ha; Los
96 Angeles: average density during the breeding season: 1.2–2.5 ind/ha; MacGregor-Fors et al.

97 2017), resulting in shorter alert distances in Mexico City when compared with Barcelona and
98 Los Angeles.

99

100 **Methods**

101 Study area

102 We conducted this study in three cities where the House Sparrow is present: Barcelona
103 (Catalonia, Spain), Los Angeles (California, United States) and Mexico City (Mexico).

104 Barcelona is located in the south-eastern region of the Iberian Peninsula (41°23'30" N, 2°10'25"
105 E; ~16 m above sea level), surrounded by the Mediterranean Sea and confined by a mountainous
106 system. It is the second most populated city in Spain (~1.6 million inhabitants), with its
107 metropolitan area housing 4.7 million residents (Demographia 2017). Los Angeles is located in
108 the south-western coast of the United States (34°3'4" N, 118°14'37" W; ~86 m above sea level)
109 (USCB 2012). The Los Angeles-Long Beach-Anaheim metropolitan area has approximately 12.1
110 million inhabitants. Finally, Mexico City is located in the Valley of Mexico, as part of the
111 Transverse Volcanic Axis (19°25'56" N, 99° 7' 59" W; ~2200 m above sea level). According to
112 the National Institute of Statistics and Geography of Mexico (INEGI 2010), the Metropolitan
113 area has ~20 million inhabitants.

114

115 Fieldwork

116 We assessed House Sparrows alert distances by walking toward House Sparrows at different
117 sectors of the studied cities during June and early July of 2016 (Barcelona: June 15–29; Los
118 Angeles: June 8–July 7; Mexico City June 1–23). In Barcelona, El Poblenou, Eixample, Sant
119 Pere, La Barceloneta, Sant Marti, Les Corts, Les Tres Torres, Sarrià-Sant Gervasi, Santa Eulàlia,

120 and Gornal; in Los Angeles, Fashion District, Flower District, Koreatown, Hollywood,
121 Inglewood, Hawthorne, and El Pueblo; in Mexico City, Pedregal de Santa Úrsula, El Caracol,
122 Villa Panamericana, and Pedregal de la Zorra. Given that after noticing our presence, House
123 Sparrows tend to hop a few meters away from its initial position and can take several repetitions
124 of such behavior before fleeing (often hopping under a car or fenced area) in some of our study
125 areas, we did not measure flight initiation distances per se, a variable that has been widely
126 studied for birds in the past (Cooke 1980; Erwin 1989; Fernández-Juricic 2000; Blumstein 2003,
127 2006; Anderson 2006). Thus, the response variable that was comparable among cities was the
128 moment in which sparrows were wary of our presence (i.e., alert distance). Upon encountering a
129 House Sparrow, one observer walked towards the targeted individual at a consistent pace (~2
130 km/h) to elicit a behavioral response. We recorded the following information when spotting a
131 House Sparrow: (1) ‘sex/age’ (i.e., adult males vs. adult females and immatures), (2) ‘flock size’
132 (number of sparrows grouped in distances <2 m with the targeted individual), and (3) distance at
133 which the individual evidently reacted to our presence approaching it (m), recorded using laser
134 rangefinders (Bushnell Yardage Pro Sport 450). We note that the sampling sites were selected
135 based previous knowledge regarding the presence of House Sparrows, as well as the accessibility
136 to conduct the surveys.

137

138 Statistical analyses

139 To test differences in the alert distances of House Sparrows at the three studied cities, we
140 performed a linear model (LM). Due to the lack of homogeneity of variance in ADs by city, we
141 log-transformed ($\log(x+1)$) ADs. We included ‘city’ (i.e., Barcelona, Los Angeles, Mexico City),
142 ‘sex/age’, and ‘flock size’ as independent variables for the LM. In order to perform contrasts

143 among the three cities, we used the ‘gmodels’ package for R (function ‘estimable’; Warnes et al.
144 2018). Based on the variables shown to be significantly related with our independent variable in
145 the LM, we performed classification and regression trees (CARTs). This method uses binary
146 recursive partitioning to identify independent variables that best explain variations in the
147 dependent variable. CARTs consider deviance, which is analogous to the residuals of sum of
148 squares in multiple regressions (used in a similar fashion to the forward procedure of
149 independent variable selection) (Crawley 2013). To run CARTs, we used the ‘rpart’ package for
150 R (Therneau et al. 2018). One of the particularities of ‘rpart’ is that it used ANOVAs to split the
151 dataset into two mutually exclusive subsets based on an identified threshold for the independent
152 variable explaining most variance of the dependent one at any given step. As the method is
153 hierarchical, sample size decreases toward the terminal, furthest down node, given by the loss of
154 deviance. Due to its procedure based in ANOVAs, ‘rpart’ is better than other ways of
155 constructing CARTs at anticipating the results of model simplification, reason why there it no
156 need to prune them. In its graphic representation, CARTs provide the mean value of the
157 dependent variable at the terminal node of all identified scenarios. Thus, CARTs are a robust
158 procedure to identify the scenarios under which the dependent variable changes in a dichotomous
159 and hierarchical manner as a result of variations of the related independent variables at each step
160 (Crawley, 2013). All statistical analyses were run in R (R Development Core Team, 2018).

161

162 **Results**

163 We gathered a total of 208 House Sparrow alert distance records in the studied cities.
164 Specifically, we recorded the response of 106 individuals in Barcelona, 50 in Los Angeles, and
165 52 in Mexico City. House Sparrow ‘flock sizes’ ranged from 1 to 10 individuals (mean \pm SD;

166 Barcelona = 1.5 ± 0.8 individuals, Los Angeles = 2.3 ± 1.9 individuals, Mexico City = 1.8 ± 1.7
167 individuals). House Sparrow alert differences were shorter for Mexico City (2.0 ± 2.7 m; max.
168 12 m), than those from the other two cities: Barcelona (8.5 ± 4.0 m; max. 20.8 m), Los Angeles
169 (7.2 ± 5.2 m; max. 23.4 m).

170 The LM showed that only ‘city’ showed a significant relationship with House Sparrow
171 ADs, while ‘sex/age’ indicated a non-significant trend ($F_{1,203} = 3.12$, $P = 0.07$), with adult males
172 being less bold than adult females and juveniles (Table 1). ADs at all three cities showed to be
173 significantly different (i.e., Barcelona–Los Angeles contrast: $t_{203} = 2.95$, $P = 0.003$; Barcelona–
174 Mexico City contrast: $t_{203} = -12.83$, $P < 0.001$; Los Angeles–Mexico City contrast: $t_{203} = 8.10$, P
175 < 0.001).

176 Considering ‘city’ and ‘sex/age’ (variables that showed a significant and a non-
177 significant trend in the LM), the CART untangled their relationship with House Sparrow alert
178 distances. Above all, it corroborated the hierarchical importance of ‘city’ in explaining recorded
179 variations in House Sparrow alert distances at the studied cities. First, alert distances were
180 shorter in Mexico City (average = 2 m), regardless of ‘sex/age’, when compared to any of the
181 four scenarios identified for Los Angeles and Barcelona. Next, the CART subsequently splits
182 Los Angeles and Barcelona, with ‘sex/age’ splitting variations in Barcelona, with adult male
183 House Sparrow exhibiting 20% larger average alert distances (9.6 m) when compared to adult
184 females and juveniles (7.9 m) (Figure 1).

185

186 **Discussion**

187 The ability to colonize and thrive in urban conditions is a key factor that facilitates the successful
188 invasion of species (Møller et al. 2015; González-Lagos and Quesada 2017). In this study, we

189 found that House Sparrows from Mexico City were significantly bolder in terms of alert
190 distances than those from Barcelona and Los Angeles (with also differed significantly among
191 them), with ‘sex/age’ representing an important variables Barcelona. Given that alert distances
192 were significantly higher in Barcelona when compared to those from Mexico City and Los
193 Angeles, our findings support our first hypothesis (relation between House Sparrow invasiveness
194 and boldness). Yet, due to the fact that alert distances were significantly lower in Mexico City
195 when contrasted to those recorded at Los Angeles, our results are also in agreement with our
196 second prediction, showing a positive association between alert distances and House sparrow
197 densities, at least in cities where it is non-native and invasive (see MacGregor-Fors et al. 2017).
198 Although it is impossible with our current dataset to determine if the sparrow boldness in Mexico
199 City is associated with its large population density in the city or if its boldness allowed it to
200 become highly successful (Nocedal 1987; Ortega-Álvarez and MacGregor-Fors 2009), our
201 results suggest that this behavior could be tied to a density-dependent process.

202 Although our results clearly show that ‘city’ was the most important variable associated
203 with shifts in House Sparrow alert distances, we also found a non-significant trend in the LM
204 with ‘sex/age’. These results were confirmed and explained by the CART, relating ‘sex/age’ only
205 with Barcelona. In this case, the analysis shows that alert distances of adult males are longer than
206 those for adult females and juveniles. Although we could not distinguish females from immature
207 males in the field, this particular finding seems counter-intuitive in the light of the behavior of
208 male House Sparrows, which have complex intra-sexual status signals and aggressive
209 interactions (e.g., bib size, wingbars; Bókony et al. 2006), and they have been shown to be more
210 resilient to certain stressors than females (Ensminger and Westneat 2012). Yet, this result could
211 also be related to differences in the dominant status of the studied House Sparrows, which have

212 been documented to adopt different behavioral strategies in urban conditions (Lendvai et al.
213 2006). Indeed, the directional survival selection of the smaller intra-sexual status signals, which
214 are positively related with boldness, has been recorded in urban passerine males in Barcelona,
215 complimenting the reasons behind male boldness in comparison to females or immatures (Senar
216 et al. 2014). Moreover, given that our study does not only consider sex, but also has an age
217 component. Age has been shown to play an important role in avian flushing distances (Dhindsa
218 and Boag 1989), and we suggest that future studies use field marks (e.g., extent of black around
219 the eye, bib size; Nakagawa and Burke 2008) and survey over specified time-windows before the
220 long breeding season of this species. In our case here, January seems to be the best option (*sensu*
221 Summers-Smith et al. 2018) to untangle the relative role of these potential drivers on House
222 Sparrow escape distances. Careful attention should be paid to local breeding periods, as this
223 species has irregular and multiple breeding events across tropical and subtropical regions
224 (Anderson 2006).

225 Regarding ‘flock size’, it was not found to be significantly related with House Sparrow
226 alert distances by the LM, and thus were not included in the CART. A previous study performed
227 in wooded parks of Madrid provided evidence that House Sparrows increase buffer distances
228 with the number of conspecifics (Fernández-Juricic et al. 2002). Yet, other studies have shown
229 that this sparrow can behave differently in heavily-urbanized sites when compared to other less-
230 developed conditions (Vincze et al. 2015). Although we did not find ‘flock size’ to relate with
231 alert distances, it is important to underline that most of our records were from birds in singles,
232 pairs, or triples (overall 1.80 ± 1.43 individuals/flock; see Results for details by city). Although
233 these numbers may seem low, they are in line with the ‘flock sizes’ reported for House Sparrows
234 in several European cities (average ‘flock size’ 1.95 individuals; Samia et al. 2017).

235 Previous studies have shown that the response of wildlife species to human approach is
236 species-specific (Blumstein 2003, 2006); yet, most studies focus on single locations, without
237 comparing populations of the same species in which their densities differ. We recognize that our
238 study considers only three cities during one field season, and therefore acknowledge the
239 limitations of the generalization of our findings. Additionally, we are aware that we only tested
240 one of the potential drivers of House Sparrow boldness, measured through alert distances; yet,
241 there are many other factors, including confounding ones, that could also be behind our findings,
242 including –but not only– habituation to humans (Møller 2008, Díaz et al. 2015), as well as
243 human density (Møller and Díaz in press; Samia et al. 2017; Vincze et al. 2015), geographical
244 differences (e.g., latitudinal behavioral responses; Díaz et al. 2013), and neuroendocrine traits
245 (Chávez-Zichinelli et al. 2010; Liebl and Martin 2012). Nevertheless, our results clearly show
246 that House Sparrow alert distances in one city located within the region where we consider it
247 native (i.e., Barcelona) were significantly larger when compared to both studied North American
248 cities. Given that alert distances were significantly larger in Los Angeles when contrasted with
249 those from Mexico City (where it is hyper-abundant; MacGregor-Fors et al. 2017), our results
250 also suggest that its boldness could be related to density-dependent processes. Based on our
251 results and others in the literature, we suggest that future studies could add evidence to or refute
252 this phenomenon by including a wider array of cities, testing hypotheses related to the
253 familiarization to humans, geographical variations, as well as the physiology of House Sparrows,
254 and including scenarios in which House Sparrows are hyper-abundant, as well as scarce, and
255 finding strategies to sex and age all of the assessed individuals.

256

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401 Table 1. LM considering relationships between ‘city,’ ‘sex/age’ (i.e., adult males vs. adult
402 females and immatures), and ‘flock size’ with House Sparrow alert distances.

403

404	Variable	F	df	P
405	‘City’	83.11	2	< 0.001
406	‘Sex/Age’	3.12	1	0.079
407	‘Flock size’	1.80	1	0.181

408 Figure 1. Classification and regression tree (CART) relating 'city' and 'sex/age' with House
 409 Sparrow alert distances in the three studied cities. Numbers displayed at the bottom of the CART
 410 represent average alert distances under the 'city' and 'sex/age' scenarios. * 'S/A' = 'sex/age'.

