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## Interspecific variation in the relationship between clutch size, laying date and intensity of urbanization in four species of hole-nesting birds

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## Abstract

The increase in size of human populations in urban and agricultural areas has resulted in considerable habitat conversion globally. Such anthropogenic areas have specific environmental characteristics, which influence the physiology, life history, and population dynamics of plants and animals. For example, the date of bud burst is advanced in urban compared to nearby natural areas. In some birds, breeding success is determined by synchrony between timing of breeding and peak food abundance. Pertinently, caterpillars are an important food source for the nestlings of many bird species, and their abundance is influenced by environmental factors such as temperature and date of bud burst. Higher temperatures and advanced date of bud burst in urban areas could advance peak caterpillar abundance and thus affect breeding phenology of birds. In order to test whether laying date advance and clutch sizes decrease with the intensity of urbanization, we analyzed the timing of breeding and clutch size in relation to intensity of urbanization as a measure of human impact in 199 nest box plots across Europe, North Africa, and the Middle East (i.e., the Western Palearctic) for four species of hole-nesters: blue tits (*Cyanistes caeruleus*), great tits (*Parus major*), collared flycatchers (*Ficedula albicollis*), and pied flycatchers (*Ficedula hypoleuca*). Meanwhile, we estimated the intensity of urbanization as the density of buildings surrounding study plots measured on orthophotographs. For the four study species, the intensity of urbanization was not correlated with laying date. Clutch size in blue and great tits does not seem affected by the intensity of urbanization, while in collared and pied flycatchers it decreased with increasing intensity of urbanization. This is the first large-scale study showing a species-specific major correlation between intensity of urbanization and the ecology of breeding. The underlying mechanisms for the relationships between life history and urbanization remain to be determined. We propose that effects of food abundance or quality, temperature, noise, pollution, or disturbance by humans may on their own or in combination affect laying date and/or clutch size.

## Introduction

The apparent dichotomy between urban and rural areas is usually used to analyze the impact of urban habitats on populations, although this definition does not consider that rural areas may also be urbanized when compared to truly natural habitats. Indeed, Pickett *et al.* (2011) defined urbanized areas as those where people live in high densities and also where infrastructures such as roads or bridges as well as buildings cover most of the surface. Urbanized areas influence climate and soil characteristics with impacts on ecosystems (Pickett *et al.* 2011). Temperatures are generally higher in cities than in neighboring rural or natural areas, phenomenon known as “heat island effect” (Escourrou 1990; Pachauri and Reisinger 2008; Stocker *et al.* 2013). These temperature increases are influenced by urban human population density (Gaston 2010; Pickett *et al.* 2011; Susca *et al.* 2011). High human population density also causes socio-political-economic pressures on ecosystems that provide services such as food, raw materials, recreational values and decontaminated water and atmosphere for human populations (Grimm *et al.* 2008; Gaston 2010; Pickett *et al.* 2011), although urban areas also support animal and plant species (Aronson *et al.* 2014).

Urbanization characteristics influence ecosystems at all levels, from individuals to communities, depending on the systematic group considered. For example, humans greatly modify plant communities in parks and gardens across cities and often urban communities are dominated by non-native plants that have lower insect populations (Pickett *et al.* 2011). A lower diversity of insects, amphibians, and reptiles occurs in urban compared to rural areas, and the abundance of domestic animals such as cats and dogs increases with human density (Bol’shakov *et al.* 2001; Gil and Brumm 2013; Johnson *et al.* 2013; Vittoz *et al.* 2013). In birds, generalists are more predominant than specialists in urban areas (Blair 1996; Devictor *et al.* 2008; Shwartz *et al.* 2008; Sorace and Gustin 2009; Huste and Boulmier 2011). Moreover, the behavior of animals, but especially also of birds, is influenced by environmental urban characteristics, for example, noisy backgrounds and/or buildings influence intra- and interspecific communication such as acoustic detection of predators and conspecifics (Brumm 2004; Barber *et al.* 2010; Snell-Rood 2012; Slabbekoorn 2013), and artificial light during night perturbs circadian and annual rhythms affecting sleep and timing of breeding (Small and Elvidge 2011; Dominoni *et al.* 2014; Fonken and Nelson 2014; Raap *et al.* 2015). Moreover, a previous quantitative review demonstrated a significant advance in laying dates in urban areas for five bird species (including great tits) and a delay for one of a total of ten avian species considered (Chamberlain *et al.* 2009).

A mismatch between phenology and suitable timing of migration or breeding may reduce individual fitness and affect population dynamics in birds (Visser *et al.* 2004, 2012). Hatching date is constrained by laying date, clutch size and incubation date (Godfray *et al.* 1991; Visser *et al.* 2004). Synchrony between individual behavior and suitable timing of breeding is determined by environmental clues (Parmesan 2006; Visser *et al.* 2006; Sih *et al.* 2011) such as temperature (Both *et al.* 2004; Charmantier *et al.* 2008; Naef-Daenzer *et al.* 2012), light (Dominoni *et al.* 2013), and date of bud burst (Visser *et al.* 2012). Moreover, global temperature increases in early spring have advanced the phenology of birds over the last few decades (Both and Visser 2001; Both *et al.* 2004; Visser *et al.* 2006; Both and te Marvelde 2007; Møller *et al.* 2010; Portier *et al.* 2012; Charmantier and Gienapp 2014; Dunn and Møller 2014). As urban areas are usually warmer than the surrounding rural areas, breeding in urban areas could be advanced by higher ambient temperatures (Escourrou 1990; Pachauri and Reisinger 2008; Stocker *et al.* 2013) and/or by artificial night light (Small and Elvidge 2011; Dominoni *et al.* 2014; Fonken and Nelson 2014), but also by more intensive feeding of birds (Stenning 1995; Robb *et al.* 2008a,b). These modifications may make urban areas become habitable to migrant birds arriving at their breeding grounds slightly earlier than nearby rural areas, which may be earlier than more natural areas (Tryjanowski *et al.* 2013; Dunn and Møller 2014).

In rural and natural habitats, egg laying of birds is delayed by cold prelaying temperatures (Charmantier *et al.* 2008; Visser *et al.* 2009; Naef-Daenzer *et al.* 2012; Schaper *et al.* 2012; Chmielewski *et al.* 2013; Vatka *et al.* 2014), at high latitudes (Mainwaring *et al.* 2012; Ruffino *et al.* 2014), or when bud burst date is delayed (Naef-Daenzer *et al.* 2012). Moreover, the both effects of latitude (Mainwaring *et al.* 2012) and bud burst (Schaper *et al.* 2011; Visser *et al.* 2012) seem to be related to temperature effects. In migratory species, laying date is mainly determined by arrival date, which in turn is advanced by global temperature increases especially in northerly populations (Walther *et al.* 2002; Both and te Marvelde 2007; Pulido 2007) even if laying date and arrival date of some migratory species are more poorly correlated than in others (Laaksonen *et al.* 2006) or in southern populations (Goodenough *et al.* 2011). Thus, variation in temperature and environmental conditions more broadly are the determinants of breeding phenology.

Previous studies of the effects of urbanization on avian life history variables have often relied on a single or a couple of populations (Hörak *et al.* 2002; Isaksson and Andersson 2007; Chamberlain *et al.* 2009; Brahmia *et al.*

2013), which does not allow for generalizations or inferences regarding spatial heterogeneity. The aim of this study was to relate breeding ecology to the intensity of local urbanization, a proxy of density and influence of humans on ecosystem, by analyzing laying dates and clutch sizes in four species of hole-nesting passerine birds, in relation to the degree of urbanization across Europe, North Africa, and the Middle East. We used hole-nesting birds as a model system because the breeding phenology is easier to follow than in open-cup nesters, and, therefore, they are routinely studied by scientists and amateurs across the Western Palearctic. This study was based on almost 200 study plots with a total of almost 80,000 reproductive events. Such extensive data are unavailable for other species of birds, but also for other organisms. These extensive data facilitated the current study. A decrease in clutch size is one option to advance hatching date (Visser et al. 2004), but could also be a response to environmental conditions in urban area as food quality, human disturbance or cat predation (Gil and Brumm 2013) or population density (Krebs 1970; Stenning et al. 1988). If intensity of urbanization did not influence laying date, we analyzed the relationship between clutch size and urban intensity with laying date as fixed factor. As ambient temperatures are higher at lower latitudes in Europe (Schönwiese and Rapp 2013), and temperature seems to be one of the main determinants of avian breeding phenology, we analyzed the interaction between latitude and intensity of urbanization on laying date. We expected a stronger impact of urbanization on tits compared to flycatchers. Indeed, flycatchers are sub-Saharan migrants that spend less time in urban areas and a laying date that is influenced by arrival date in northern populations, while laying date of tits is mainly determined by local conditions at the breeding sites (Pearson and Lack 1992; Both et al. 2006). Finally, laying date may also vary with habitat structure (Van Balen 1973; Mänd et al. 2005; Arriero et al. 2006; Mizuta 2006) or the presence of predators (Lank and Ydenberg 2003; Sergio et al. 2007). Larger and deeper nest boxes (Mertens 1977; Van Balen 1984; Summers and Taylor 1996) and concrete boxes (O'Connor 1978) offer better thermal isolation and better protection from predation. Thus, the influence of dominant habitat, nest floor surface, and nest box material on laying date and clutch size were also analyzed.

## Material and Methods

### Study species

Blue tits (*Cyanistes caeruleus*), great tits (*Parus major*), pied flycatchers (*Ficedula hypoleuca*), and collared flycatchers (*Ficedula albicollis*) are all small insectivorous

passerine birds that breed commonly in nest boxes in large parts of Europe. The two species of tits are residents or partial short-distance migrants depending on their population (Nowakowski and Vähätalo 2003), while the two flycatchers species are both migratory and spend the winter months in sub-Saharan West Africa.

### Data

Annual mean breeding dates, clutch sizes, and sample sizes of first clutches derive from an exhaustive attempt to obtain information from populations across Europe, North Africa, and the Middle East (Fig. S1; Møller et al. 2014). We used mean laying date per population and year of (1) 101 study populations of blue tits, with a total of 1127 study years for laying date, and 1124 study years for clutch size; (2) 138 study populations of great tits, 1439 study years for laying date, and 1436 for clutch size; (3) 66 study populations of collared flycatchers, with a total of 592 study years for both parameters; and (4) 23 populations of pied flycatchers, with a total of 259 study years. Information on latitude, longitude, altitude, mean study year, species, dominant breeding habitat (coniferous, deciduous, evergreen, or mixed forest), nest box floor area, and nest box material (wood or concrete) for all study plots were provided by scientists or reported in a previous publication (Møller et al. 2014). Borders and numbers of nest boxes depended on study plot as determined by researchers who monitored the populations.

The increase in the density of buildings is known to correlate with the increase in the total number of individual birds (Hedblom and Söderström 2010), advanced laying date (Shustack and Rodewald 2010), and increased breeding success (Ryder et al. 2010; Hedblom and Söderström 2012). All study plots were classified by each researcher who followed a given population as either rural or urban without single criterion (see Supporting information). This dichotomy is commonly used, but inadequate when quantifying human influence because rural areas include agricultural, and natural habitats and urban areas include parks and gardens with mature trees. Therefore, we recorded an estimate of the density of buildings using information from ArcGIS Earthstar Geographics for each of the study plots ( $N = 199$ ). An index of the "Intensity of urbanization" was obtained by dividing the number of buildings by the area of the study site (see next paragraph) followed by log-transformation of density of buildings adding a constant of one to avoid values of zero and to normalize the data. We counted the number of roofs of each building as the number of roofs with one color and one direction, while L-shaped buildings were counted as two roofs. We used the density of buildings and not the cover by roads or buildings because these



descriptive variables are strongly positively correlated (Shustack and Rodewald 2010). The percentage of built-up area within 1-km circles was strongly positively correlated with the number of roofs ( $F_{1,32} = 81.24$ ,  $P < 0.01$ ), and previous studies have shown that it is the density of buildings that is correlated with population density (Brumm 2004; Barber *et al.* 2010; Snell-Rood 2012; Slabekoor 2013). Study plot coordinates were in the form 12.12345°N, 12.12345°E for 160 study plots and in the form 12.12°N, 12.12°E for 39 study plots where the scientist was inaccessible either due to retirement or death, and it was thus impossible to provide more precise coordinates (see Table S1). Analyses were repeated without these imprecise coordinates for study plots, but we found qualitatively similar results.

To estimate the effect of measurement scale, we analyzed the intensity of urbanization in subsamples of 34 of 199 study plots (all study plots classified as urban [ $N = 14$ ] and 20 randomly selected from 185 plots classified as rural plots by scientists [see Table S1]) within a radius of 200, 500, and 1000 m from the centre of each study plot, by visually counting the number of buildings on digital orthophotographs (Shustack and Rodewald 2010). The density of buildings was measured at the scale that allowed identification of different roofs, depending on the orthophotographs available. We used a radius of 200 and 500 m, respectively, because they were similar to the size of most study plots and 1000 m to validate the method at the level of study plots. Use of a buffer circular area around study plots provided conservative estimates of intensity of urbanization among study plots. However, the intensity of urbanization at the three distances was highly repeatable (200–500 m,  $F_{33,34} = 8.20$ ,  $P < 0.01$ , intraclass correlation coefficient = 0.78 (Lessells and Boag 1987) and 200–1000 m,  $F_{33,34} = 3.30$ ,  $P < 0.01$ , intraclass correlation coefficient = 0.54). Therefore, we only used an estimate of the intensity of urbanization ( $\log[\text{number of building/area of study} + 1]$ ) for a radius of 200 m in the subsequent analyses; that is, an intensity of urbanization was recorded in each of the study plots in 2015 ( $N = 199$ ). Our indicator of intensity of urbanization was on average 59 buildings/km<sup>2</sup> (SE = 4, range 0–1305 buildings/km<sup>2</sup>,  $N = 199$ ). We found a strong positive relationship between the binomial score of urbanization provided by scientists and intensity of urbanization near the nest box plots for all study plots (Student *t*-test:  $t_{df} = -6.26_{14,19}$ ,  $P < 0.0001$ , Mean  $\pm$  SE log-transformed index =  $0.47 \pm 0.06$  and  $2.15 \pm 0.26$  for rural ( $N = 185$ ) and urban ( $N = 14$ ) areas, respectively, see Fig. S1A). Moreover, intensity of urbanization was negatively related to CORINE land cover code (Kendall rank order test:  $\tau = -0.46$ ,  $t_{df} = -7.11_{189}$ ,  $P < 0.0001$ ,  $N = 191$ , see Fig. S1B). CORINE land cover code assessed the land

cover in classes (agricultural areas, artificial surfaces or forests areas), with values decreasing with degree of anthropogenization of areas. These codes were available for most areas of Europe, but not North Africa and Middle East. Intensity of urbanization in agricultural areas was intermediate between that in urban and natural sites.

For some species, latitude is correlated with laying date (Mainwaring *et al.* 2012; Ruffino *et al.* 2014), and to take this into account, geographic coordinates of study plots were used in the models. Latitude and longitude of the study plots were positively correlated (Pearson  $r = 0.48$ ,  $t_{197} = 7.64$ ,  $P < 0.01$ , Fig. S2), although not causing problems of collinearity (based on correlograms; Dormann *et al.* 2007). The interaction between longitude and latitude and quadratic terms for latitude and longitude were entered in models to account for nonlinear relationships and spatial autocorrelation (Legendre 1993). There was no autocorrelation in model residuals (Moran test; Dormann *et al.* 2007).

Collared flycatchers only used wooden nest boxes, while the two species of flycatchers were absent from evergreen habitats.

## Statistical analyses

All statistical analyses were performed in R v. 3.2.0 (R Core Team 2015). Explanatory variables were correlated, but coefficients were small (see Table S2). We used linear mixed models and backward elimination of factors using Akaike's information criterion (AIC) to select the best predictive model to explain variation in laying date (package nlme, function lme, method REML and package car, function Anova, type III). The initial model included the three-way interaction (latitude  $\times$  species  $\times$  intensity of urbanization), two-way interactions (latitude  $\times$  species, species  $\times$  intensity of urbanization, latitude  $\times$  intensity of urbanization and latitude  $\times$  longitude), study plot as random factor and intensity of urbanization, log-transformed altitude, latitude, latitude squared, longitude, longitude squared, nest floor surface, material of nest box and dominant habitat as fixed factors. As the three-way interaction was significant (see Table S3), models were subsequently developed and performed for each of the four species separately.

In the second part, for species for which the interaction between latitude and intensity of urbanization and the main effect of intensity of urbanization were not significant, the initial model to explain variation in clutch size included two-way interactions (latitude  $\times$  intensity of urbanization and latitude  $\times$  longitude), study plot as random factor and intensity of urbanization, laying date, log-transformed altitude, latitude, latitude squared, longitude, longitude squared, nest floor surface, material of nest box,

and dominant habitat as fixed factors. Altitude was log-transformed to avoid residuals of models that deviated from normal distributions. As discussed in detail above, we used linear mixed models and backward elimination of factors using AIC (package nlme, function lme, method REML and package car, function Anova, type III).

Several nests were followed in each plot and each year (blue tits: from 1 to 154 nests, mean  $\pm$  SE =  $17 \pm 22$  nests per year and per plot/great tits: from 1 to 210 nests, mean  $\pm$  SE =  $20 \pm 24$  nests per year and per plot/collared flycatchers: from 1 to 159 nests, mean  $\pm$  SE =  $26 \pm 19$  nests per year and per plot/pied flycatchers: from 1 to 189 nests, mean  $\pm$  SE =  $50 \pm 35$  nests per year and per plot). The number of nests per year and per study plot was used for weighting each data point, thereby assuring that each observation contributed to the models relative to the level of sampling (Draper and Smith 1998; Kutner *et al.* 2004). Likelihood ratio tests (LRT) were manually calculated for the random effect of study plot. No residuals of final models deviated from normal distributions.

## Results

### Laying date

Box plots of laying dates for the four species across a gradient of “intensity of urbanization” are shown in Figure 1. For all four species, the interaction between latitude and intensity of urbanization, the main effect of intensity of urbanization, and altitude were not significant, while laying date advanced significantly over years (Table 1).

In blue tits, laying date varied with nest box floor area and study plot coordinates, and differed significantly among habitats (Table 1). In great tits, laying date varied with study plot coordinates, and it differed among habitats (Table 1). In collared flycatchers, laying date was correlated with study plot coordinates, nest box floor area, and habitat (Table 1). In pied flycatchers, laying date varied with study plot coordinates and was earlier in wooden nest boxes compared to concrete boxes (Table 1).

### Clutch size

Box plots of clutch size for the four species across a gradient of “intensity of urbanization” are shown in Figure 2. For all four species, the interaction between latitude and intensity of urbanization and altitude were not significant, and clutch sizes did not differ between habitats but decreased with laying date (Table 2).

In blue tits, clutch size did not vary significantly with the intensity of urbanization, but decreased across years, varied with nest box floor area, nest box material and

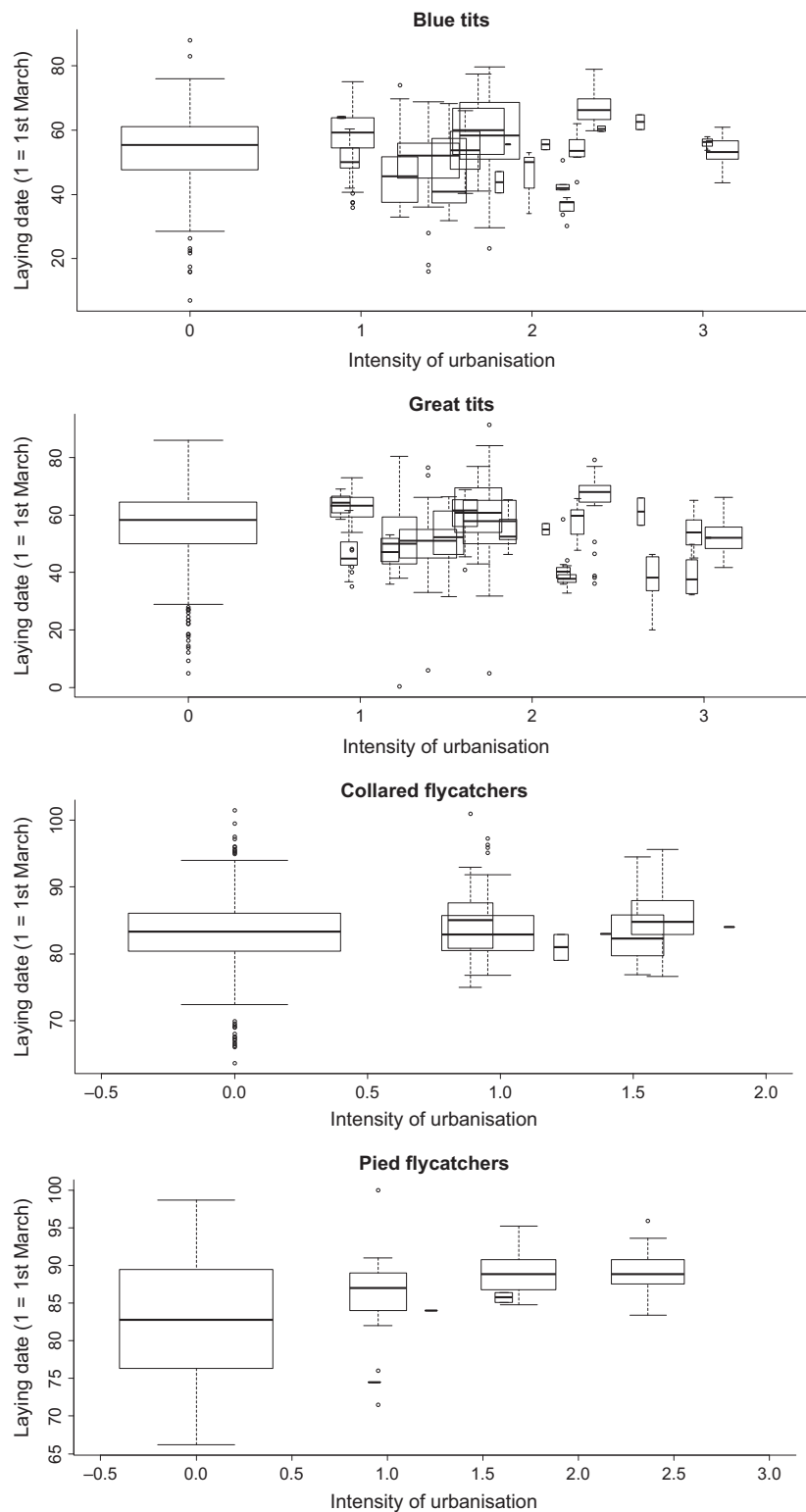
study plot coordinates (Table 2). In great tits, clutch size did not vary significantly with intensity of urbanization, but decreased across years and varied with study plot (Table 2). In collared flycatchers, clutch size decreased with intensity of urbanization and increased across years (Table 2). In pied flycatchers, clutch size decreased with intensity of urbanization, varied with study plot coordinates and was larger in wooden than in concrete nest boxes (Table 2).

## Discussion

We analyzed the breeding ecology of four species of passerine birds in nest boxes in relation to the intensity of urbanization across Europe, North Africa, and the Middle East. Nest box characteristics, habitat, and geographic location were included in the models to account for potentially confounding environmental effects other than that of intensity of urbanization. To our knowledge, this is the first large-scale study of the relationship between intensity of urbanization and avian breeding ecology. The intensity of urbanization was not correlated with laying date in the four species, while clutch sizes decreased with increasing intensity of urbanization in both collared and pied flycatchers.

In all four species, our large-scale analysis confirmed correlations between laying date, clutch size, and various environmental factors which have previously been demonstrated in single-specific studies (Van Balen 1973, 1984; Mertens 1977; O'Connor 1978; Pearson and Lack 1992; Summers and Taylor 1996; Both and Visser 2001; Ahola *et al.* 2004; Both *et al.* 2004, 2006; Mänd *et al.* 2005; Arriero *et al.* 2006; Mizuta 2006; Both and te Marvelde 2007; Charmantier *et al.* 2008; Magi *et al.* 2009; Sisask *et al.* 2010; Mainwaring *et al.* 2012; Chmielewski *et al.* 2013; Charmantier and Gienapp 2014; Møller *et al.* 2014; Ruffino *et al.* 2014; Vatka *et al.* 2014). Thus, we focus the remainder of the discussion on the correlation between intensity of urbanization and breeding ecology.

Urbanization is an ongoing process that has intensified over time and differs among countries. Indeed, temporal changes in urbanization varied between 0.1% per year in the Netherlands measured in 1992 (WRR 1992) to more than 2% per year in France between 2000 and 2010 (Clanché and Rascol 2011). With these differences in rate of urbanization, it was difficult to take temporal change in urbanization into account. In order to verify the validity of our measure of intensity of urbanization among years, we only analyzed data collected after 2000. However, we still found qualitatively similar results (analyses not shown). Thus, the use of a unique intensity of urbanization for a specific year seemed not to be an issue for the analysis of the correlation between urbanization on



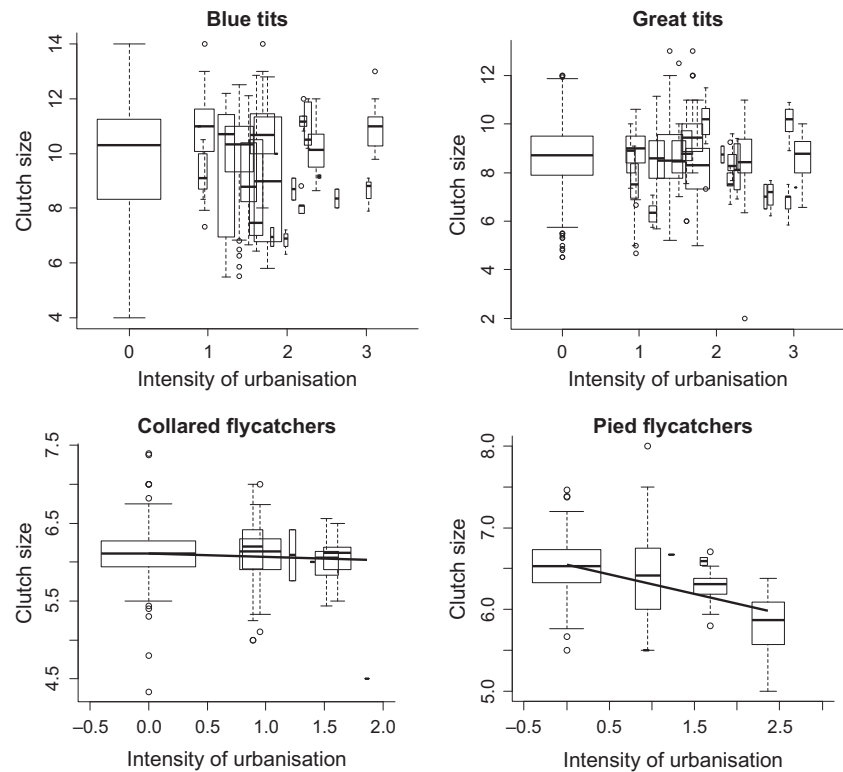
**Figure 1.** Box plots of laying date in relation to intensity of urbanization in four passerine bird species in Europe, North Africa, and the Middle East. Box plots show medians, quartiles, 5- and 95-percentiles, and extreme values. Width of box plots reflects sample size (study populations/ total number of years: 100/1125, 138/1439, 66/592, and 23/259 for blue tit, great tit, collared flycatcher, and pied flycatcher, respectively). Intensity of urbanization was estimated as the local density of buildings per km<sup>2</sup> and log-transformed.



**Table 1.** Results of linear models investigating variation in laying date as a function of two-way interactions (latitude  $\times$  intensity of urbanization and latitude  $\times$  longitude), study plot (random factor), intensity of urbanization, altitude (log-transformed), latitude, longitude and longitude squared, year, nest floor area, nest box material, and dominant habitat (fixed factors).

Species	Blue tit	Great tit	Collared flycatcher	Pied flycatcher								
Study populations	101	138	66	23								
Total number of study years	1127	1439	592	259								
Final (initial) model AIC	7837.31 (7840.73)	10,513.13 (10,526.64)	3875.78 (3879.74)	1764.44 (1795.32)								
Intensity of urbanization	$F_{df}$ 0.78, 98	$P$ 0.39	Estimate $\pm$ SE -1.16 $\pm$ 1.32	$F_{df}$ 0.21, 132	$P$ 0.65	Estimate $\pm$ SE 3.01 $\pm$ 6.61	$F_{df}$ 0.15, 60	$P$ 0.70	Estimate $\pm$ SE 269.66 $\pm$ 690.06	$F_{df}$ 0.27, 18	$P$ 0.60	Estimate $\pm$ SE 14.86 $\pm$ 28.47
Latitude	9.87 <sub>1,1014</sub>	<b>&lt;0.01</b>	-9.64 $\pm$ 3.07	2.12 <sub>1,134</sub>	0.15	-3.93 $\pm$ 2.70	0.96 <sub>1,60</sub>	0.33	8.27 $\pm$ 8.44	192.14 <sub>1,234</sub>	<b>&lt;0.01</b>	1.74 $\pm$ 0.13
Latitude <sup>2</sup>	11.40 <sub>1,1014</sub>	<b>&lt;0.01</b>	0.11 $\pm$ 0.03	2.42 <sub>1,134</sub>	0.12	0.04 $\pm$ 0.03	0.21 <sub>1,56</sub>	0.65	-0.39 $\pm$ 0.84	0.04 <sub>1,227</sub>	0.84	-0.02 $\pm$ 0.10
Longitude	6.94 <sub>1,1014</sub>	<b>&lt;0.01</b>	3.61 $\pm$ 1.37	2.53 <sub>1,134</sub>	0.11	-0.89 $\pm$ 0.56	0.70 <sub>1,60</sub>	0.40	18.71 $\pm$ 22.35	0.01 <sub>1,227</sub>	0.92	0.21 $\pm$ 3.46
Longitude <sup>2</sup>	5.70 <sub>1,1014</sub>	<b>0.02</b>	0.06 $\pm$ 0.03	23.50 <sub>1,134</sub>	<b>&lt;0.01</b>	-0.05 $\pm$ 0.01	0.10 <sub>1,56</sub>	0.75	-0.49 $\pm$ 1.52	0.01 <sub>1,227</sub>	0.92	-0.01 $\pm$ 0.02
Year	161.99 <sub>1,14014</sub>	<b>&lt;0.01</b>	-0.19 $\pm$ 0.02	36.00 <sub>1,1292</sub>	<b>&lt;0.01</b>	-0.09 $\pm$ 0.01	34.83 <sub>1,524</sub>	<b>&lt;0.01</b>	-0.15 $\pm$ 0.03	24.51 <sub>1,234</sub>	<b>&lt;0.01</b>	-0.15 $\pm$ 0.03
Altitude (log)	0.11 <sub>1,1014</sub>	0.74	0.66 $\pm$ 1.95	1.48 <sub>1,1292</sub>	0.29	1.47 $\pm$ 1.40	0.74 <sub>1,524</sub>	0.39	-3.24 $\pm$ 3.78	1.55 <sub>1,18</sub>	0.21	-2.05 $\pm$ 1.65
Nest floor area	6.37 <sub>1,1014</sub>	<b>0.01</b>	-0.05 $\pm$ 0.02	0.30 <sub>1,1291</sub>	0.59	0.01 $\pm$ 0.01	0.31 <sub>1,523</sub>	0.58	-0.04 $\pm$ 0.07	1.54 <sub>1,227</sub>	0.22	-0.02 $\pm$ 0.01
Nest box material	1.25 <sub>1,1014</sub>	0.26		1.16 <sub>1,1292</sub>	0.28		-	-		4.72 <sub>1,21</sub>	<b>0.03</b>	
Wood			-3.59 $\pm$ 3.21			2.48 $\pm$ 2.31			-			4.45 $\pm$ 2.05
Dominant habitat	60.56 <sub>3,1014</sub>	<b>&lt;0.01</b>		12.65 <sub>1,1292</sub>	<b>&lt;0.01</b>		0.04 <sub>2,56</sub>	0.98		0.06 <sub>2,227</sub>	0.97	
Deciduous			-13.75 $\pm$ 2.00			-1.44 $\pm$ 1.98			-0.33 $\pm$ 1.60			-0.31 $\pm$ 2.21
Evergreen			-10.26 $\pm$ 2.08			5.61 $\pm$ 2.75			-			-
Mixed			-4.36 $\pm$ 1.01			-2.56 $\pm$ 1.26			-0.41 $\pm$ 3.50			-0.33 $\pm$ 1.35
Intensity of urbanization $\times$ latitude	0.10 <sub>1,1013</sub>	0.76	0.06 $\pm$ 0.19	0.25 <sub>1,1291</sub>	0.62	-0.07 $\pm$ 0.13	0.15 <sub>1,60</sub>	0.70	-4.73 $\pm$ 12.10	0.25 <sub>1,18</sub>	0.62	-0.23 $\pm$ 0.47
Latitude $\times$ longitude	7.40 <sub>1,1014</sub>	<b>&lt;0.01</b>	-0.09 $\pm$ 0.03	13.22 <sub>1,134</sub>	<b>&lt;0.01</b>	0.04 $\pm$ 0.01	0.74 <sub>1,60</sub>	0.39	-0.38 $\pm$ 0.44	0.01 <sub>1,227</sub>	0.97	-0.01 $\pm$ 0.07
Study plot	LRT: -28.41	0.50		LRT: 404.70	<b>&lt;0.01</b>		LRT: 523.46	<b>&lt;0.01</b>		LRT: 351.304	<b>&lt;0.01</b>	

Initial values of variables that were not retained in the final models are presented in italics, and significant *P*-values in the final models are shown in bold on a gray background.



**Figure 2.** Box plots of clutch size in relation to intensity of urbanization in four passerine bird species in Europe, North Africa, and the Middle East. Box plots show medians, quartiles, 5- and 95-percentiles, and extreme values. Width of box plot reflects sample size (study populations/total number of years: 100/1122, 138/1436, 66/592, and 23/259 for blue tit, great tit, collared flycatcher, and pied flycatcher, respectively). Lines are the linear regression. Intensity of urbanization was estimated as the local density of buildings per km<sup>2</sup> and log-transformed.

breeding phenology. Indeed, even if European cities did not grow at the same speed, a highly urbanized city in mid-century is also likely to be equally highly urbanized today (EEA 2015).

Laying date was not related to the intensity of urbanization in any of the four species. The lack of significant relationship between the intensity of urbanization and laying date in all four species could be due to lack of sensitivity to urbanization or due to the proxy, the intensity of urbanization, used to quantify the degree of human impact on the environment. According to the relationship between CORINE land cover code and the intensity of urbanization, the index measured was related to anthropogenization of areas although radius could still be too small for some borders of monitored study plots. Although we studied local urbanization of study plots, it is still possible that human impact affects the environment at larger scales (Bol'shakov *et al.* 2001; Pickett *et al.* 2011; Gil and Brumm 2013; Johnson *et al.* 2013; Vittoz *et al.* 2013). The findings could be affected by the lack of highly urbanized plots, because the plots sampled in our study did not cover all variation in the intensity of urbanization in all European countries. Data were limited by availability of boxes differing in extent of urbanization, although we consider that this is not a serious issue in the present study because rural plots included natural plots as forests and agricultural or industrial plots where

people also live. Nevertheless, we have shown that laying dates of the four species were not related to the intensity of local urbanization and lack of data does not seem to affect this result.

Collared and pied flycatchers showed a significant negative relationship between the intensity of urbanization and clutch size. Migratory status could be the decisive factor for the decrease in clutch size in flycatchers and the absence of such a difference in tits. The lack of a significant effect in tits could also be due to differences in thermal capacity as blue and great tits live at more variable latitudes than collared and pied flycatchers (Fig. S3; Svensson 1992; Del Hoyo *et al.* 2007) and hence display a larger range of temperature tolerance. This is the first time that a negative relationship has been shown between clutch size and intensity of urbanization in the two long-distance migratory flycatcher species (Both *et al.* 2004, 2006; Laaksonen *et al.* 2006; Mizuta 2006; Pulido 2007; Sisask *et al.* 2010; Massa *et al.* 2011; Smallegange *et al.* 2011). More studies are needed to understand the underlying mechanism of intensity of urbanization on clutch size, and it is even possible that a reduction in clutch size was due to a combination of local conditions (Pearson and Lack 1992; Both *et al.* 2006) determined by ambient temperature (Burrows *et al.* 2011; Stocker *et al.* 2013), artificial night light (Small and Elvidge 2011; Dominoni *et al.* 2014; Fonken and Nelson 2014), food availability

**Table 2.** Results of linear models investigating variation in clutch size as a function of two-way interactions (latitude  $\times$  intensity of urbanization and latitude  $\times$  longitude), study plot (random factor), intensity of urbanization, laying date, altitude (log-transformed), latitude, latitude squared, longitude and longitude squared, year, nest floor area, nest box material, and dominant habitat (fixed factors).

Species	Blue tit	Great tit	Collared flycatcher	Pied flycatcher
Study populations	101	138	66	23
Total number of study years	1124	1436	592	259
Final (initial) model AIC	3911.17 (3958.37)	5611.25 (5645.29)	956.86 (993.03)	640.50 (712.91)
Intensity of urbanization	$F_{df}$ $0.02_{,1,98}$	$P$ 0.88 $Estimate \pm SE$ $-0.13 \pm 0.88$ $F_{df}$ $0.05_{,1,32}$	$P$ 0.82 $Estimate \pm SE$ $0.17 \pm 0.75$ $F_{df}$ 9.66 <sub>1,64</sub>	$P$ <b>0.04</b> $Estimate \pm SE$ $-0.24 \pm 0.08$ $F_{df}$ 4.18 <sub>1,20</sub>
Laying date	46.15 <sub>1,1018</sub>	<b>&lt;0.01</b> $-0.033 \pm 0.005$	<b>&lt;0.01</b> $-0.039 \pm 0.005$	<b>&lt;0.01</b> $-8.65 \pm 4.73$
Latitude	8.90 <sub>1,1018</sub>	<b>&lt;0.01</b> $0.70 \pm 0.23$	<b>&lt;0.01</b> $0.82 \pm 0.30$	<b>&lt;0.01</b> $9.00 \pm 6.99$
Latitude <sup>2</sup>	4.68 <sub>1,1018</sub>	<b>0.03</b> $-0.05 \pm 0.002$	<b>&lt;0.01</b> $-0.008 \pm 0.003$	<b>0.96</b> $-0.13 \pm 0.10$
Longitude	1.37 <sub>1,1010</sub>	0.24 $0.17 \pm 0.15$	0.24 $-0.08 \pm 0.06$	0.30 $-5.10 \pm 5.76$
Longitude <sup>2</sup>	2.75 <sub>1,1010</sub>	0.10 $0.004 \pm 0.003$	0.28 $-0.01 \pm 0.01$	0.35 $-0.26 \pm 0.18$
Year	29.04 <sub>1,14018</sub>	<b>&lt;0.01</b> $-0.015 \pm 0.003$	<b>&lt;0.01</b> $-0.021 \pm 0.003$	0.22 $0.009 \pm 0.002$
Altitude (log)	1.68 <sub>1,1010</sub>	0.19 $-0.26 \pm 0.20$	0.22 $0.21 \pm 0.18$	0.77 $-0.20 \pm 0.11$
Nest floor area	12.64 <sub>1,1018</sub>	<b>&lt;0.01</b> $-0.010 \pm 0.003$	0.44 $0.01 \pm 0.01$	0.12 $0.01 \pm 0.01$
Nest box material	7.44 <sub>1,1018</sub>	<b>&lt;0.01</b> $0.93_{,1,288}$	0.34 $0.93_{,1,288}$	<b>&lt;0.01</b> $15.49_{,1,20}$
Wood		0.83 $\pm$ 0.30	0.28 $\pm$ 0.29	–
Dominant habitat	4.55 <sub>3,1010</sub>	0.21	0.46 $0.01_{,2,56}$	0.53 $1.27_{,2,226}$
Deciduous		0.43 $\pm$ 0.25	0.16 $\pm$ 0.26	0.01 $\pm$ 0.23
Evergreen		0.21 $\pm$ 0.30	–0.22 $\pm$ 0.36	–
Mixed		0.33 $\pm$ 0.18	0.22 $\pm$ 0.22	–0.01 $\pm$ 0.42
Intensity of urbanization	0.01 <sub>1,1010</sub>	0.96 $-0.01 \pm 0.02$	0.76 $-0.01 \pm 0.01$	0.50 $0.06 \pm 1.56$
× latitude				
Latitude	1.83 <sub>1,1010</sub>	0.18 $-0.005 \pm 0.004$	<b>&lt;0.01</b> $0.003 \pm 0.001$	0.29 $0.30 \pm 0.23$
× longitude				
Study plot	LRT: 790.62	LRT: 1384.18	LRT: 754.05	LRT: 496.76
		<b>&lt;0.01</b>	<b>&lt;0.01</b>	<b>&lt;0.01</b>

Initial values of variables that were not retained in the final models are presented in *italics*, and significant *P*-values in the final models are shown in **bold** on a gray background.

(Stenning 1995; Robb et al. 2008a,b; Saggese et al. 2011), avian population density (Krebs 1970; Stenning et al. 1988), nest predation by cats (Zanette et al. 2011), or vandalism (Brahmia et al. 2013).

In conclusion, in this first large-scale study of life history traits and intensity of urbanization we showed a complex species-specific major relationship between intensity of urbanization and breeding. The underlying mechanisms for the relationships between life history and intensity of urbanization remain to be determined. However, we propose that effects of food abundance or quality, avian population density, temperature, noise, pollution or disturbance by humans may on their own or in combination affect laying date and/or clutch size. Experiments could compare main and interactive effects of bird feeding by humans (Chamberlain et al. 2009) and temperature increases on advances in breeding date in neighboring urban and rural habitats.

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## Conflict of Interest

None declared.

## References

- Ahola, M., T. Laaksonen, K. Sippola, T. Eeva, K. Rainio, and E. Lehikoinen. 2004. Variation in climate warming along the migration route uncouples arrival and breeding dates. *Glob. Change Biol.* 10:1610–1617.
- Aronson, M. F. J., F. A. La Sorte, C. H. Nilon, M. Katti, M. A. Goddard, C. A. Lepczyk, et al. 2014. A global analysis of the impacts of urbanization on bird and plant diversity reveals key anthropogenic drivers. *Proc. R. Soc. B Biol. Sci.* 281:20133330.
- Arriero, E., J. J. Sanz, and M. Romero-Pujante. 2006. Habitat structure in Mediterranean deciduous oak forests in relation to reproductive success in the blue tit *Parus caeruleus*. *Bird Study* 53:12–19.
- Barber, J. R., K. R. Crooks, and K. M. Fristrup. 2010. The costs of chronic noise exposure for terrestrial organisms. *Trends Ecol. Evol.* 25:180–189.
- Blair, R. B. 1996. Land use and avian species diversity along an urban gradient. *Ecol. Appl.* 6:506–519.
- Bol'shakov, V. N., O. A. Pyastolova, and V. L. Vershinin. 2001. Specific features of the formation of animal species communities in technogenic and urbanized landscapes. *Russ. J. Ecol.* 32:315–325.
- Both, C., and L. te Marvelde. 2007. Climate change and timing of avian breeding and migration throughout Europe. *Clim. Res.* 35:93–105.
- Both, C., and M. E. Visser. 2001. Adjustment to climate change is constrained by arrival date in a long-distance migrant bird. *Nature* 411:296–298.
- Both, C., A. V. Artemyev, B. Blaauw, R. J. Cowie, A. J. Dekhuijzen, T. Eeva, et al. 2004. Large-scale geographical variation confirms that climate change causes birds to lay earlier. *Proc. R. Soc. B Biol. Sci.* 271:1657–1662.
- Both, C., J. J. Sanz, A. V. Artemyev, B. Blaauw, R. J. Cowie, A. J. Dekhuijzen, et al. 2006. Pied Flycatchers *Ficedula hypoleuca* travelling from Africa to breed in Europe: differential effects of winter and migration conditions on breeding date. *Ardea* 94:511–525.
- Brahmia, Z., R. Scheifler, N. Crini, S. Maas, P. Giraudoux, and S. Benyacoub. 2013. Breeding performance of blue tits (*Cyanistes caeruleus ultramarinus*) in relation to lead pollution and nest failure rates in rural, intermediate, and urban sites in Algeria. *Environ. Pollut.* 174:171–178.
- Brumm, H. 2004. The impact of environmental noise on song amplitude in a territorial bird. *J. Anim. Ecol.* 73:434–440.
- Burrows, M. T., D. S. Schoeman, L. B. Buckley, P. Moore, E. S. Poloczanska, K. M. Brander, et al. 2011. The pace of shifting climate in marine and terrestrial ecosystems. *Science* 334:652–655.
- Chamberlain, D. E., A. R. Cannon, M. P. Toms, D. I. Leech, B. J. Hatchwell, and K. J. Gaston. 2009. Avian productivity in urban landscapes: a review and meta-analysis. *The Ibis* 151:1–18.
- Charmantier, A., and P. Gienapp. 2014. Climate change and timing of avian breeding and migration: evolutionary versus plastic changes. *Evol. Appl.* 7:15–28.
- Charmantier, A., R. H. McCleery, L. R. Cole, C. Perrins, L. E. B. Kruuk, and B. C. Sheldon. 2008. Adaptive phenotypic plasticity in response to climate change in a wild bird population. *Science* 320:800–803.
- Chmielewski, F. M., K. Blumel, C. Scherbaum-Heberer, B. Koppmann-Rumpf, and K. H. Schmidt. 2013. A model approach to project the start of egg laying of great tit (*Parus major* L.) in response to climate change. *Int. J. Biometeorol.* 57:287–297.

- Clanché, F., and O. Rascol. 2011. Le découpage en unités urbaines de 2010. *Insee Première* 1364:1–4.
- Del Hoyo, J., A. Elliott, and D. A. Christie. 2007. *Handbook of the birds of the world*. Lynx Editions, Barcelona, Spain.
- Devictor, V., R. Julliard, and F. Jiguet. 2008. Distribution of specialist and generalist species along spatial gradients of habitat disturbance and fragmentation. *Oikos* 117: 507–514.
- Dominoni, D., M. Quetting, and J. Partecke. 2013. Artificial light at night advances avian reproductive physiology. *Proc. R. Soc. Lond. B Biol. Sci.* 280:20123017.
- Dominoni, D. M., E. O. Carmona-Wagner, M. Hofmann, B. Kranstauber, and J. Partecke. 2014. Individual-based measurements of light intensity provide new insights into the effects of artificial light at night on daily rhythms of urban-dwelling songbirds. *J. Anim. Ecol.* 83:681–692.
- Dormann, C. F., J. M. McPherson, M. B. Araújo, R. Bivand, J. Bolliger, G. Carl, et al. 2007. Methods to account for spatial autocorrelation in the analysis of species distributional data: a review. *Ecography* 30:609–628.
- Draper, N. R., and H. Smith. 1998. *Applied regression analysis*. Probability and statistics. 3rd ed. John Wiley & Sons, New York.
- Dunn, P. O., and A. P. Møller. 2014. Changes in breeding phenology and population size of birds. *J. Anim. Ecol.* 83:729–739.
- EEA. 2015. *European environment: state and outlook 2015: assessment of global megatrends*. European Environment Agency, Copenhagen, Denmark.
- Escourrou, G. 1990. La spécificité du climat de l'agglomération parisienne/The unique character of the Paris urban area's climate. *Rev. Géogr. Lyon* 65:85–89.
- Fonken, L. K., and R. J. Nelson. 2014. The effects of light at night on circadian clocks and metabolism. *Endocr. Rev.* 35:648–670.
- Gaston, K. J. 2010. *Urban ecology*. Cambridge Univ. Press, Cambridge, U.K.
- Gil, D., and H. Brumm. 2013. *Avian urban ecology*. Oxford Univ. Press, Oxford, U.K.
- Godfray, H. C. J., L. Partridge, and P. H. Harvey. 1991. Clutch size. *Annu. Rev. Ecol. Syst.* 22:409–429.
- Goodenough, A., A. Hart, and S. Elliot. 2011. What prevents phenological adjustment to climate change in migrant bird species? Evidence against the “arrival constraint” hypothesis. *Int. J. Biometeorol.* 55:97–102.
- Grimm, N. B., S. H. Faeth, N. E. Golubiewski, C. L. Redman, J. G. Wu, X. M. Bai, et al. 2008. Global change and the ecology of cities. *Science* 319:756–760.
- Hedblom, M., and B. Söderström. 2010. Landscape effects on birds in urban woodlands: an analysis of 34 Swedish cities. *J. Biogeogr.* 37:1302–1316.
- Hedblom, M., and B. Söderström. 2012. Effects of urban matrix on reproductive performance of great tit (*Parus major*) in urban woodlands. *Urban Ecosyst.* 15:167–180.
- Hörak, P., P. F. Surai, and A. P. Møller. 2002. Fat-soluble antioxidants in the eggs of great tits *Parus major* in relation to breeding habitat and laying sequence. *Avian Sci.* 2:123–130.
- Huste, A., and T. Boulinier. 2011. Determinants of bird community composition on patches in the suburbs of Paris, France. *Biol. Conserv.* 144:243–252.
- Isaksson, C., and S. Andersson. 2007. Carotenoid diet and nestling provisioning in urban and rural great tits *Parus major*. *J. Avian Biol.* 38:564–572.
- Johnson, P. T. J., J. T. Hoverman, V. J. McKenzie, A. R. Blaustein, and K. L. D. Richgels. 2013. Urbanization and wetland communities: applying metacommunity theory to understand the local and landscape effects. *J. Appl. Ecol.* 50:34–42.
- Krebs, J. R. 1970. Regulation of numbers in the Great tit (Aves: Passeriformes). *J. Zool.* 162:317–333.
- Kutner, M. H., C. J. Nachtsheim, J. Neter, and W. Li. 2004. *Applied linear statistical models*. 5th revised ed. McGraw-Hill Irwin, New York.
- Laaksonen, T., M. Ahola, T. Eeva, R. A. Väisänen, and E. Lehikoinen. 2006. Climate change, migratory connectivity and changes in laying date and clutch size of the pied flycatcher. *Oikos* 114:277–290.
- Lank, D. B., and R. C. Ydenberg. 2003. Death and danger at migratory stopovers: problems with “predation risk”. *J. Avian Biol.* 34:225–228.
- Legendre, P. 1993. Spatial autocorrelation: trouble or new paradigm? *Ecology* 74:1659–1673.
- Lessells, C. M., and P. T. Boag. 1987. Unrepeatable repeatabilities: a common mistake. *Auk* 104:116–121.
- Magi, M., R. Mänd, H. Tamm, E. Sisask, P. Kilgas, and V. Tilgar. 2009. Low reproductive success of great tits in the preferred habitat: a role of food availability. *Ecoscience* 16:145–157.
- Mainwaring, M. C., I. R. Hartley, S. Bearhop, K. Brulez, C. R. du Feu, G. Murphy, et al. 2012. Latitudinal variation in blue tit and great tit nest characteristics indicates environmental adjustment. *J. Biogeogr.* 39:1669–1677.
- Mänd, R., V. Tilgar, A. Lõhmus, and A. Leivits. 2005. Providing nest boxes for hole-nesting birds – does habitat matter? *Biodivers. Conserv.* 14:1823–1840.
- Massa, B., C. A. Cusimano, B. Margagliotta, and R. Galici. 2011. Reproductive characteristics and differential response to seasonal temperatures of blue and great tits (*Cyanistes caeruleus* & *Parus major*) in three neighbouring mediterranean habitats. *Rev. Écol.(Terre Vie)* 66:157–172.
- Mertens, J. A. L. 1977. Thermal conditions for successful breeding in great tits (*Parus major* L.) II. Thermal properties of nests and nestboxes and their implications for the range of temperature tolerance of great tit broods. *Oecologia* 28:31–56.
- Mizuta, T. 2006. Habitat difference in food abundance possibly causes within-population variation in the timing of



- breeding in the Madagascar paradise flycatcher (*Terpsiphone mutata*). *Can. J. Zool.* 84:817–822.
- Møller, A. P., W. Fiedler, and P. Berthold. 2010. Effects of climate change on birds. Oxford Univ. Press, Oxford, U.K.
- Møller, A. P., F. Adriaensen, A. Artemyev, J. Banbura, E. Barba, C. Biard, et al. 2014. Clutch-size variation in Western Palaearctic secondary hole-nesting passerine birds in relation to nest box design. *Methods Ecol. Evol.* 5:353–362.
- Naef-Daenzer, B., J. Luterbacher, M. Nuber, T. Rutishauser, and W. Winkel. 2012. Cascading climate effects and related ecological consequences during past centuries. *Clim. Past* 8:1527–1540.
- Nowakowski, J. K., and A. V. Vähätalo. 2003. Is the great tit *Parus major* an irruptive migrant in north-east Europe? *Ardea* 91:231–243.
- O'Connor, R. J. 1978. Nest-box insulation and the timing of laying in the Wytham woods population of great tits *Parus major*. *The Ibis* 120:534–537.
- Pachauri, R. K., and A. Reisinger. 2008. *Bilan 2007 des changements climatiques: Rapport de synthèse*. Rapport d'évaluation du Groupe d'experts intergouvernemental sur l'évolution du climat GIEC, Genève, Switzerland.
- Parmesan, C. 2006. Ecological and evolutionary responses to recent climate change. *Annu. Rev. Ecol. Evol. Syst.* 37:637–669.
- Pearson, D. J., and P. C. Lack. 1992. Migration patterns and habitat use by passerine and near-passerine migrant birds in eastern Africa. *The Ibis* 134:89–98.
- Pickett, S. T. A., M. L. Cadenasso, J. M. Grove, C. G. Boone, P. M. Groffman, E. Irwin, et al. 2011. Urban ecological systems: scientific foundations and a decade of progress. *J. Environ. Manage.* 92:331–362.
- Porlier, M., A. Charmantier, P. Bourgault, P. Perret, J. Blondel, and D. Garant. 2012. Variation in phenotypic plasticity and selection patterns in blue tit breeding time: between- and within-population comparisons. *J. Anim. Ecol.* 81:1041–1051.
- Pulido, F. 2007. Phenotypic changes in spring arrival: evolution, phenotypic plasticity, effects of weather and condition. *Clim. Res.* 35:5–23.
- R Core Team. 2015. R: a language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria.
- Raap, T., R. Pinxten, and M. Eens. 2015. Light pollution disrupts sleep in free-living animals. *Sci. Rep.* 5:13557.
- Robb, G. N., R. A. McDonald, D. E. Chamberlain, and S. Bearhop. 2008a. Food for thought: supplementary feeding as a driver of ecological change in avian populations. *Front. Ecol. Environ.* 6:476–484.
- Robb, G. N., R. A. McDonald, D. E. Chamberlain, S. J. Reynolds, T. J. E. Harrison, and S. Bearhop. 2008b. Winter feeding of birds increases productivity in the subsequent breeding season. *Biol. Lett.* 4:220–223.
- Ruffino, L., P. Salo, E. Koivisto, P. B. Banks, and E. Korpimäki. 2014. Reproductive responses of birds to experimental food supplementation: a meta-analysis. *Front. Zool.* 11:1–13.
- Ryder, T. B., R. Reitsma, B. Evans, and P. P. Marra. 2010. Quantifying avian nest survival along an urbanization gradient using citizen- and scientist-generated data. *Ecol. Appl.* 20:419–426.
- Saggese, K., F. Korner-Nievergelt, T. Slagsvold, and V. Amrhein. 2011. Wild bird feeding delays start of dawn singing in the great tit. *Anim. Behav.* 81:361–365.
- Schaper, S. V., C. Rueda, P. J. Sharp, A. Dawson, and M. E. Visser. 2011. Spring phenology does not affect timing of reproduction in the great tit (*Parus major*). *J. Exp. Biol.* 214:3664–3671.
- Schaper, S. V., A. Dawson, P. J. Sharp, P. Gienapp, S. P. Caro, and M. E. Visser. 2012. Increasing temperature, not mean temperature, is a cue for avian timing of reproduction. *Am. Nat.* 179:E55–E69.
- Schönwiese, C. D., and J. Rapp. 2013. Climate trend atlas of Europe based on observations 1891–1990. Springer Science & Business Media, Springer, Dordrecht, the Netherlands.
- Sergio, F., L. Marchesi, P. Pedrini, and V. Penteriani. 2007. Coexistence of a generalist owl with its intraguild predator: distance-sensitive or habitat-mediated avoidance? *Anim. Behav.* 74:1607–1616.
- Shustack, D. P., and A. D. Rodewald. 2010. Attenuated nesting season of the Acadian flycatcher (*Empidonax virens*) in urban forests. *Auk* 127:421–429.
- Shwartz, A., S. Shirley, and S. Kark. 2008. How do habitat variability and management regime shape the spatial heterogeneity of birds within a large Mediterranean urban park? *Landscape Urban Plan.* 84:219–229.
- Sih, A., M. C. O. Ferrari, and D. J. Harris. 2011. Evolution and behavioural responses to human-induced rapid environmental change. *Evol. Appl.* 4:367–387.
- Sisask, E., R. Mänd, M. Magi, and V. Tilgar. 2010. Parental provisioning behaviour in pied flycatchers *Ficedula hypoleuca* is well adjusted to local conditions in a mosaic of deciduous and coniferous habitat. *Bird Study* 57:447–457.
- Slabbekoorn, H. 2013. Songs of the city: noise-dependent spectral plasticity in the acoustic phenotype of urban birds. *Anim. Behav.* 85:1089–1099.
- Small, C., and C. D. Elvidge. 2011. Mapping decadal change in anthropogenic night light. Pp. 353–358 in A. Stein, E. Pebesma, G. Heuvelink, eds. *Spatial statistics 2011: mapping global change*. Elsevier Science, Amsterdam, the Netherlands.
- Smallegange, I. M., J. van der Meer, and W. Fiedler. 2011. Population dynamics of three songbird species in a nestbox population in Central Europe show effects of density, climate and competitive interactions. *The Ibis* 153:806–817.
- Snell-Rood, E. C. 2012. The effect of climate on acoustic signals: does atmospheric sound absorption matter for bird song and bat echolocation? *J. Acoust. Soc. Am.* 131:1650–1658.

- Sorace, A., and M. Gustin. 2009. Distribution of generalist and specialist predators along urban gradients. *Landscape Urban Plan.* 90:111–118.
- Stenning, M. 1995. Hatching asynchrony in a population of blue tits *Parus caeruleus*. [PhD thesis], University of Sussex, Sussex.
- Stenning, M. J., P. H. Harvey, and B. Campbell. 1988. Searching for density-dependent regulation in a population of pied flycatchers *Ficedula hypoleuca* Pallas. *J. Anim. Ecol.* 57: 307–317.
- Stocker, T. F., D. Qin, G.-K. Plattner, M. Tignor, S. K. Allen, J. Boschung, et al. 2013. *Climate change 2013: the physical science basis*. Contribution of working group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change. Cambridge, U.K. and New York.
- Summers, R. W., and W. G. Taylor. 1996. Use by tits of nest boxes of different designs in pinewoods. *Bird Study* 43:138–141.
- Susca, T., S. R. Gaffin, and G. R. Dell'Osso. 2011. Positive effects of vegetation: urban heat island and green roofs. *Environ. Pollut.* 159:2119–2126.
- Svensson, L. 1992. Identification guide to European passerines. British Trust for Ornithology, Norfolk, United Kingdom.
- Tryjanowski, P., T. H. Sparks, S. Kuzniak, P. Czechowski, and L. Jerzak. 2013. Bird migration advances more strongly in urban environments. *PLoS One* 8:e63482.
- Van Balen, J. H. 1973. A comparative study of the breeding ecology of the great tit *Parus major* in different habitats. *Ardea* 61:1–93.
- Van Balen, J. 1984. The relationship between nest-box size, occupation and breeding parameters of the great tit *Parus major* and some other hole-nesting species. *Ardea* 72:163–175.
- Vatka, E., S. Rytönen, and M. Orell. 2014. Does the temporal mismatch hypothesis match in boreal populations? *Oecologia* 176:595–605.
- Visser, M. E., C. Both, M. M. Lambrechts, A. Møller, W. Fiedler, and P. Berthold. 2004. Global climate change leads to mistimed avian reproduction. *Adv. Ecol. Res.* 00:89–110. Academic Press.
- Visser, M. E., L. J. M. Holleman, and P. Gienapp. 2006. Shifts in caterpillar biomass phenology due to climate change and its impact on the breeding biology of an insectivorous bird. *Oecologia* 147:164–172.
- Visser, M. E., L. J. M. Holleman, and S. P. Caro. 2009. Temperature has a causal effect on avian timing of reproduction. *Proc. R. Soc. B Biol. Sci.* 276:2323–2331.
- Visser, M. E., L. te Marvelde, and M. E. Lof. 2012. Adaptive phenological mismatches of birds and their food in a warming world. *J. Ornithol.* 153:S75–S84.
- Vittoz, P., D. Cherix, Y. Gonseth, V. Lubini, R. Maggini, N. Zbinden, et al. 2013. Climate change impacts on biodiversity in Switzerland: a review. *J. Nat. Conserv.* 21:154–162.
- Walther, G. R., E. Post, P. Convey, A. Menzel, C. Parmesan, T. J. C. Beebee, et al. 2002. Ecological responses to recent climate change. *Nature* 416:389–395.
- WRR. 1992. Ground for choices. Four perspectives for the rural areas in the European Community. Netherlands Scientific Council for Government Policy, The Hague, the Netherlands.
- Zanette, L. Y., A. F. White, M. C. Allen, and M. Clinchy. 2011. Perceived predation risk reduces the number of offspring songbirds produce per year. *Science* 334:1398–1401.

## Supporting Information

Additional Supporting Information may be found online in the supporting information tab for this article:

**Figure S1.** Intensity of urbanisation according to (A) classification by scientists. Box plots show medians, quartiles, 5- and 95-percentiles, and extreme values, and (B) CORINE land cover code (red = discontinuous urban, purple = industrial or commercial units, pink = green urban sites, brown = arable land and rice field, orange = agriculture lands, green = forest and natural field and blue = inland marshes).

**Figure S2.** Distribution of study plots across Europe, North Africa and the Middle East.

**Figure S3.** Box plots of latitude of study plots in four passerine birds in Europe, North Africa and the Middle East.

**Table S1.** Summary data for study plots. See Material and methods for definitions.

**Table S2.** Correlation matrix of explanatory variables.

**Table S3.** Mixed linear model investigating laying date in four passerines species (CF: Collared Flycatcher, GT: Great tit and PF: Pied Flycatcher) as a function of habitat characteristics (intensity of urbanisation, latitude, longitude squared, longitude, longitude squared, altitude (log-transformed), and dominant habitat), nest box characteristics (nest floor surface and nest box material) and year as fixed effects, with study plot as a random factor.