## Interactive effects of fearfulness and geographical location on bird population trends

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## ABSTRACT

Animal populations are currently under pressure from multiple factors that include human land use and climate change. They may compensate for such effects by reducing, either by habituation or by natural selection, the distance at which they flee from humans (i.e., flight initiation distance, FID), and this adaptation may improve their population trends. We analyzed population trends of common breeding birds in relation to FID and geographical location (latitude, longitude, and marginality of the breeding distribution) across European countries from Finland in the north to Spain in the south, while also considering other potential predictors of trends like farmland habitat, migration, body size and brain size. We found evidence of farmland, migratory and smaller-sized species showing stronger population declines. In contrast, there was no significant effect of relative brain size on population trends. We did not find evidence for main effects of FID and geographical location on trends after accounting for confounding and interactive effects; instead, FID and location interacted to generate complex spatial patterns of population trends. Trends were more positive for fearful populations northwards, westwards and (marginally) towards the centre of distribution areas, and more negative for fearless populations toward the south, east, and the margins of distribution ranges. These findings suggest that it is important to consider differences in population trends among countries, but also interaction effects among factors, because such interactions can enhance or compensate for negative effects of other factors on population trends.

Key-words: breeding birds, flight initiation distance, latitude, longitude, marginality of distribution.

## INTRODUCTION

Human disturbance of wild organisms is a common cause of concern in a world with a rapidly increasing human population (Wong and Candolin 2012; Ehrlich and Ehrlich 2013). Such effects of disturbance include release of stress hormones (Wingfield and Ramenofsky 1999), increased metabolic rate (Belanger and Bédard 1990), reduction in foraging activity (Madsen 1998a; 1998b), displacement from preferred foraging and roosting sites and changes in diurnal rhythms (Madsen and Fox 1995) and non-lethal effects of predation (Abrams 1991). These factors on their own and in combination may have effects on the condition of animals and hence on their reproduction and survival prospects. A common behavioral measure of proneness to disturbance by humans and animals alike is the flight initiation distance (FID): The distance at which an animal takes flight when approached by a potential predator (Cooper and Blumstein 2015). Because all animals continuously have to weigh the risk of falling prey to a predator by fleeing too late when approached against the benefits of staying put and hence continuing to feed and/or rest, FID constitutes an instantaneous measure of this individual trade-off. Cooke (1980) noticed that urban birds had much shorter flight distances than rural populations of the same species, and that this difference depended on body size, the difference being larger in small species with high metabolism. This change in behavior between urban and rural habitats allowed birds to coexist with humans even at high human population densities, which are a cause of frequent disturbance. Parallel latitudinal trends in FID and raptor abundance in paired urban and rural sites suggest that birds, besides responding to human presence, also adjust their behavior in response to natural levels of disturbance by predators (see Díaz et al. 2013 and references therein).

It has been noticed that human disturbance at seabird colonies linked to escape behavior and FID could result in altered habitat use and reduced reproductive performance (Burger 1981; Burger and Gochfeld 1981). Therefore, FID can be a useful tool in conservation including assessment of levels of disturbance and susceptibility to disturbance (Madsen 1995; 1998a; 1998b;

Weston et al. 2012). The population consequences of FIDs can be investigated by relating population trends to FID (Møller 2008). We should expect species with long FIDs for their body size to show declining population trends because such species should be more prone to get disturbed by humans. Among 56 species of birds, FID accounted for $33 \%$ of the variance in population trends in Denmark, with effect sizes ranging from 0.36 to 0.58 in different analyses. Therefore, species with long FIDs for their body size had declining populations while species with short FIDs had increasing populations even when controlling statistically for potentially confounding effects. However, a study on population trends in the UK in relation to predictors that included FID recorded in Denmark did not find significant relationship between FID and population trends (Thaxter et al. 2010). This raises the question whether population trends and FID should originate from the same geographic location to make analyses meaningful.

Many national and international monitoring programs tally population trends of organisms as diverse as birds, mammals, butterflies and bumblebees. In particular, birds have been popular targets for monitoring since the 1960's in many countries in Europe, and population trends based on European continentwide monitoring have been published since 1980 (European Bird Census Council, http://www.ebcc.info/index.php?ID=509). According to these data, while many species have increased in distribution and abundance, a majority, at least in specific habitats such as farmland, have shown a clear decline. Although humans either directly or indirectly play a major role in determining long-term population trends of birds in Europe (Reif 2013), the underlying mechanisms remain poorly understood. In addition, population trends vary across the distribution range of species. Cuervo and Møller (2013) found stronger increases in northern populations and greater fluctuations in marginal populations, somewhat expected from influences of global warming on climatic niches (Hampe and Petit 2005), and Donald, Green and Heath (2001) and Reif et al. (2011) showed longitudinally varying trends. Reif et al. (2011) also showed an interesting difference in the effect of relative brain size on trends at both sides of
the iron curtain, consistent with the differences in land-use intensity across Europe. These intriguing and varying patterns, and the need to optimize conservation priorities, mean that there are good reasons to investigate patterns of population trends at different spatial scales in an attempt to elucidate the underlying mechanisms, including the potential effects of FID.

The objectives of this study were to test whether population trends were related to FID, and whether these influences varied across the European continent. If spatial changes in FID could partially compensate for the main effects of factors of global change on trends, we predicted significant interactions between FID and latitude, longitude and marginality on trends. We also tested whether previously established predictors of population trends such as farmland habitat, migration distance, body mass or brain mass affected the relationship between population trend and FID. Overall, elucidating geographical variation in the relationships between trends and recent responses of organisms to changes in the level of human activities will help us to understand our impact on wild populations of animals and eventually to reduce such impacts.

## METHODS

We recorded FID for a total of 159 species during the breeding seasons 20092010 at nine locations from eight countries along a wide latitudinal gradient across Europe, from Finland in the north to Spain in the south, by using a standard procedure developed by Blumstein (2006). These data are reported in Díaz et al. (2013). In brief, we walked at ordinary walking speed towards a bird recording the distance from the bird when we started walking, the distance at which the birds initiated escape, and the bird's height in the vegetation. This information was used to estimate FID. In order to account for the height at which individuals were perched, FID was calculated as the Euclidean distance between the approaching human and the focal bird (which equals the square-root of the sum of the squared flight distance and the squared height in the
vegetation). Observers wore neutrally colored clothes and behaved as normal pedestrians. FID was measured by a number of trained observers and therefore data were pooled for analysis. We used the FID estimates for rural populations in each location, which consisted of paired rural and urban sites (Díaz et al. 2013), because the population size estimates used to assess trends for each country are mostly based on data coming from non-urban populations (Cuervo and Møller 2013). Data for the two Spanish sites were averaged to obtain a single country-level estimate.

Population trends for breeding birds in all European countries for which we had information on FID (Finland, Norway, Denmark, Poland, Czech Republic, Hungary, France and Spain) were obtained from Cuervo and Møller (2013). Available population size estimates for each bird species and country were regressed on years, and the slope of this regression was used as a proxy for population trend. We used time series of 7-27 years gathered until 2004-2008 depending on countries and species (see Cuervo and Møller 2013 for details and a full discussion of the quality of trend estimates).

Latitude and longitude for each country were estimated as the coordinates of the mid-point between the northernmost and the southernmost, and between the easternmost and the westernmost, mainland points of every country, excluding islands except for Denmark. Latitude and longitude for each country were considered the latitude and longitude for all bird populations in that particular country regardless of the actual distribution of every species within the country. Marginality of each bird population was estimated by comparing two distances (in degrees): L is the distance between the population (i.e., the country) latitude and the northernmost or the southernmost (the one that resulted in a shorter distance) limits of the breeding distribution range of the species. L was set to zero in the few cases in which the country latitude index was more southern than the southernmost limit of the species range or more northern than the northernmost limit of the species range. C is the distance between the population latitude and the latitude of the mid-point between the northernmost
and the southernmost limits of the breeding distribution range of the species. Marginality was computed as $\log _{10}(\mathrm{C}+1)-\log _{10}(\mathrm{~L}+1)$, with positive values representing marginal populations (the distance to the range centre was larger than the distance to the nearest limit) and negative values central populations (the distance to the range centre was smaller than the distance to the nearest limit). These values were transformed by adding the absolute value of the most negative number and dividing by the largest value resulting from the previous addition, to ensure that marginality estimates ranged from 0 (central population) to 1 (marginal population; see Cuervo and Møller 2013 for details).

Bird population trends have previously been shown to be systematically affected by body size, migration distance, farmland habitat and relative brain size (reviews in Møller 2008; Møller, Rubolini and Lehikoinen 2008; Reif 2013). We extracted information on mean body mass of adult birds of each species from Cramp and Perrins (1977-1994). Migration distances (mean of the northernmost and the southernmost latitudes of the breeding distribution range minus the corresponding mean for the wintering distribution range) were taken from Møller, Rubolini and Lehikoinen (2008). Farmland habitat was coded as 1 (species depending on arable and/or mixed farmland) or 0 (species depending on other habitat types) following Appendix 2 in Tucker and Evans (1997). Relative brain sizes were the residuals of a log-log phylogenetically corrected regression of brain mass on body mass based on a sample of 567 bird species (Møller, 2008); brain mass data were obtained from Garamszegi, Møller and Erritzøe (2002), Iwaniuk and Nelson (2002), Galván and Møller (2011) and Møller and Erritzøe (2014).

We $\log _{10}$-transformed FID, population trend and migration distance before analyses. Within-species repeatability of FID and trends across Europe was computed following Lessells and Boag (1987), and differences between them and the null hypothesis of zero repeatability were tested following Becker (1984). Significant repeatabilities imply statistical dependence of estimates for the same species in different countries, a fact that will bias results based on
phylogenetically-structured databases (Garamszegi and Møller 2010). As species occupy a variable number of study locations and countries (Díaz et al. 2013; Cuervo and Møller 2013), geographical trends could be partly due to phylogenetic effects. To control for such relationships we used phylogenetic generalized least square regression (PGLS) models implemented in R (Díaz et al. 2013). After estimating the phylogenetic scaling parameter lambda ( $\lambda$ ), we calculated the phylogenetically corrected partial correlations between the variables of interest. Different populations of the same species were considered as polytomies with a constant small genetic distance of $10^{-10}$ between them. We used the R script and the edited phylogeny supplied as Supplementary Files S1 and S2 in Díaz et al. (2013), but using the function pglm3.3.r instead of the pglm3.1.r to fit type III (orthogonal) models. We used the phylogeny reported in Thuiller et al. (2011). The dependent variable was the population trend, confounding variables were farmland habitat, migration distance, body size and relative brain size, and predictors FID, latitude, longitude, marginality, and the first-order interaction between FID and geographical variables. Predictors were computed from the corresponding input variables $\left(\log _{10}\right.$ FID and geographical variables) by standardizing them (i.e., by subtracting sample means and dividing by standard deviations), in order to allow direct comparison of effect sizes (Pearson's product-moment correlation coefficients computed from $P$ values of t-tests according to Lipsey and Wilson 2001) and to make main effects biologically interpretable even when involved in interactions (Schielzeth 2010).

## RESULTS

We collected data on mean FID and on recent population trends from 338 populations of 129 bird species. Data on farmland habitat, body size and migration distance were available for all of them, while there were no data on brain size for 9 species (Appendix A). Both FID and trends were significantly repeatable within species ( $F_{1,209}=3.08, P<0.001$ and $F_{1,209}=1.45, P=0.009$, respectively). FID was significantly more repeatable than population trends ( $r=$
$0.45 \pm 0.04(\mathrm{SD}) v s . r=0.15 \pm 0.05 ; t_{338}=4.0, P<0.001$; Becker 1984); in other words, geographical variation within species was larger for population trends than for mean fearfulness as reflected by FID.

Log-transformed population trends were significantly related to $\log _{10}$ FID $\left(F_{1,337}=7.96, P=0.005, r^{2}=0.02\right)$, but not to latitude $\left(F_{1,337}=0.00, P=0.967\right.$, $\left.r^{2}=0.00\right)$, longitude $\left(F_{1,337}=0.40, P=0.530, r^{2}=0.00\right)$ or marginality $\left(F_{1,337}=\right.$ $0.62, P=0.432, r^{2}=0.00$ ) when predictor effects were analyzed one by one. The relationship with FID vanished, however, after correcting for significant effects of farmland habitat, migration distance and body mass (effect sizes for these three confounding variables ranged from 0.14 to 0.16 ), while also accounting for phylogenetic effects (Table 1). Trends were more negative for farmland birds, long-distance migrants and smaller species (Table 1). Relative brain size showed no significant effects on population trends, which did not show significant geographical trends either (Table 1). However, FID showed significant interactive effects with latitude and longitude, and marginallysignificant interactive effects with marginality, with effect sizes ranging from 0.10 to 0.13 (Table 1, Fig. 1). FID-trend relationships were more positive northwards, westwards and (marginally) towards the centre of distribution areas (Table 1, Fig. 1). These interactions implied that trends were more negative for fearless populations toward the south, east, and the margins of distribution ranges.

## DISCUSSION

Many different factors have been proposed to account for population trends of birds (reviewed in Reif 2013). These variables range from migration and the perils of living under different climate regimes (Hjort and Lindholm 1978; Baillie and Peach 1992; Sanderson et al. 2006; Reif 2013), relative brain mass that facilitates the ability to cope with changing environments (Shultz et al. 2005; Møller, Rubolini and Lehikoinen 2008; Reif et al. 2011), thermal range and hence the ability to cope with changing climatic conditions (Jiguet et al.

2007, 2010), the number of broods with species producing more broods doing better (Julliard, Jiguet and Couvet 2004), and body mass with large sized species with smaller total populations having negative population trends (Bennett and Owens 2002).

Geographical variation in trends within breeding ranges of species are also expected due to geographical changes in the suitability of environmental conditions (the niche variation hypothesis; Brown 1984), in the intensity of global change drivers (Hampe and Petit 2005; Reif et al. 2011; Tryjanowski et al. 2011) or in both (Díaz et al. 1998). It has been suggested that population responses of birds to environmental gradients may be highly species-specific, even precluding broad generalizations (Taper, Böhning-Gaese and Brown 1995); however, Cuervo and Møller (2013) have recently shown that changes in population size of breeding birds in Europe are the strongest at the margins of the breeding distribution, but are particularly negative at the southern-most range margins, where increasing temperatures may render environmental conditions for maintenance of viable populations the most difficult. Climate change has affected the distribution of many species, and range margins have on average moved pole-wards (Chen et al. 2011), and recent work has shown finegrained effects of climate change on local population trends (Jiguet et al. 2010). Longitudinal variation due to differences in land-use intensity between Western and Eastern Europe have also been documented, especially for farmland birds (Donald, Green and Heath 2001). However, we did not find evidence for direct effects of these variables after accounting for effects of third variables and their interactions. Reif et al. (2011) suggested that longitudinal effects of the iron curtain dividing industrialized Western Europe from more extensive land use in Eastern Europe interacted with relative brain mass to account for spatial heterogeneity in population trends. Here we found no evidence of an effect of relative brain mass on population trends contrary to previous reports (Shultz et al. 2005; Thaxter et al. 2010; Reif et al. 2011). We hypothesize that these differences among studies may arise from the inclusion of different predictors
and their interactions, but also from inclusion of multiple countries that differ in significant predictors of population trends. Studies such as this, encompassing the widest ranges of variation of relevant variables available even at the expense of lower precision within ranges, are thus essential to detect non-linear and interactive relationships of geographically-varying conditions on local abundance and trends (e.g. Jiguet et al. 2010; Concepción et al. 2012).

Bird species breeding on farmland displayed the steepest declines. This is probably a consequence of agriculture having become ever more industrialized and intensified and thereby disproportionately negatively affecting farmland specialists (Fuller et al. 1995; Chamberlain et al. 2000; Møller, Rubolini and Lehikoinen 2008; Reif 2013). Here we found evidence consistent with this general trend, with farmland species showing more negative population trends than non-farmland birds. Migration has been predicted to affect population trends because migrants are affected negatively by land-use and climate change in their breeding range, during migration and in their winter quarters (Hjort and Lindholm 1978; Baillie and Peach 1992; Sanderson et al. 2006; Møller, Rubolini and Lehikoinen 2008; Reif 2013). Here we found a negative effect of migration distance on population trends, when accounting for the effects of the remaining variables.

We hypothesized that population trends would be negatively related to FID, as reported by Møller (2008) for European birds. Most recent work indicates that FID can be considered a general measure of the willingness of animals to be involved in risky activities such as foraging and courtship under perceived risky conditions (reviewed in Cooper and Blumstein 2014, 2015). Such willingness to take risks would depend on levels of risk (abundance and identity of predators and other sources of risk, such as humans), but also on potential fitness benefits (ie. it will be worth taking more risks if the expected fitness consequence of the reward is larger, as under food shortage or timelimited conditions), after accounting for species- and population-specific proneness to risk-taking associated with phylogeny, urban habitat or life-history
traits such as body size or migratory behavior (Díaz et al. 2013). We found an overall main effect of FID in this study, which however vanished when considering interactive effects with geographical location. This fact suggests that the observed geographical variation in trends would in fact be the net result of complex interactions between spatial variations in many factors proposed to drive population trends (Reif 2013), as well as on the varying effects of risktaking behaviors on trends. Our results showed that fearfulness of bird populations (i.e., long FIDs) enhanced population trends where such trends were already less negative, as in northern European populations (Cuervo and Møller 2013), or where land use intensity is higher, as in western European countries (Tryjanowski et al. 2011), but these relationships reversed at more stressful extremes of spatial gradients, such as southern and marginal locations. We interpret these interactions as implying that we cannot assess predictors by considering solely their main effects. We are unaware of any previous studies investigating such interaction effects as predictors of population trends.

In conclusion, we have analyzed for the first time how geographical patterns of population trends of birds in Europe, as related to natural and manmade geographical variation in environmental factors such as climate, predation risk and land use, interact with a measure of the tolerance of birds to human disturbance. Overall we found that proneness to risk-taking as estimated by short FIDs enhanced population resilience to disturbance in a changing world, as more tolerant individuals will reduce the costs associated with escape behaviors (Cooper and Blumstein 2014). In contrast, bird species and populations less tolerant of frequent disturbance, by humans or wild and domestic predators, would perform worse, especially at the southern- and eastern-most edges of breeding distributions. Further studies including fine-grained estimates of FID, trends and secondary influences on them (eg. Jiguet et al. 2010) carried out over wide geographical gradients would be needed to ascertain whether these patterns were due to geographical variations in risks, fitness benefits of risk-taking, or both.

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## APPENDIX A. SUPPLEMENTARY MATERIAL

Information on bird species, country, population trend, mean FID (m), migration distance ( ${ }^{\circ}$ latitude), body mass (g), latitude and longitude of the population/country, marginality of the population within the species breeding range, relative brain size and farmland habitat. Relative brain size is residuals from a log-log phylogenetically corrected regression of brain mass on body mass. See Material and methods for sources and details. Nomenclature and basic phylogeny follows Thuiller et al. (2011).

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Fig. 1. Latitudinal and longitudinal variation in standardized regression coefficients ( $\mathrm{B} \pm \mathrm{SE}$ ) between population trends of European birds (residuals from a multiple regression between log-transformed trends, migration distance, body mass, relative brain mass, farmland habitat, marginality, and either longitude or latitude) and fearfulness (flight initiation distance, FID, logtransformed). Lines are best-fit regressions.



| Source | estimate (SE) | $t$ | $P$ | Effect size |
| :--- | :---: | :---: | :---: | :---: |
| Farmland | $-0.008(0.003)$ | -2.50 | 0.013 | 0.14 |
| Migration distance | $-0.006(0.002)$ | -2.71 | 0.007 | 0.15 |
| Body mass | $0.018(0.006)$ | 2.93 | 0.004 | 0.16 |
| Relative brain size | $-0.014(0.010)$ | -1.46 | 0.147 | 0.08 |
| FID | $0.000(0.002)$ | 0.16 | 0.876 | 0.01 |
| Latitude | $-0.002(0.002)$ | -1.07 | 0.286 | 0.06 |
| Longitude | $0.003(0.002)$ | 1.50 | 0.134 | 0.08 |
| Marginality | $0.001(0.002)$ | 0.38 | 0.706 | 0.02 |
| FID x Latitude | $0.005(0.002)$ | 2.29 | 0.023 | 0.13 |
| FID x Longitude | $-0.004(0.002)$ | -2.27 | 0.024 | 0.12 |
| FID x Marginality | $-0.003(0.002)$ | -1.76 | 0.079 | 0.10 |

Table 1. Relationships between population trends of European birds (response variable) and geographical location (latitude, longitude and marginality) and fearfulness (flight initiation distance, FID), after accounting for effects of farmland habitat, migration distance, body mass and relative brain size on trends and correcting for the effect of the phylogenetic structure of the data set, that was, however, not significant $\left(\lambda=0.000, \chi^{2}=-0.012, P=1.000\right)$. The full model (no removal of non-significant terms was done, as recommended by Forstmeier and Schielzeth 2011) had the statistics $F=4.73$, d.f. $=12,329$, adjusted $r^{2}=$ $0.11, P<0.0001$. Effect sizes are Pearson's product-moment correlation coefficients.

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## APPENDIX A. SUPPLEMENTARY MATERIAL

Information on bird species, country, population trend, mean FID (m), migration distance ( ${ }^{\circ}$ latitude), body mass (g), latitude and longitude of the population/country, marginality of the population within the species breeding range, relative brain size and farmland habitat. Relative brain size is residuals from a log-log phylogenetically corrected regression of brain mass on body mass. See Material and methods for sources and details. Nomenclature and basic phylogeny follows Thuiller et al. (2011).

| Species | Country | Popul. trend | FID (m) | Migration distance ( ${ }^{\circ}$ latitude) | Body mass <br> (g) | Latitude | Longitude | Marginality | $\begin{gathered} \hline \text { Rel. brain } \\ \text { size } \\ \hline \end{gathered}$ | Farmland |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Accipiter nisus | Denmark | 0.002 | 36.14 | 12.79 | 204.0 | 56.16 | 10.38 | 0.585 | 0.464 | 0 |
| Acrocephalus palustris | Czech Rep. | -0.010 | 14.14 | 66.84 | 12.0 | 49.80 | 15.48 | 0.599 | -0.281 | 0 |
| Acrocephalus palustris | Denmark | 0.012 | 8.76 | 66.84 | 12.0 | 56.16 | 10.38 | 0.683 | -0.281 | 0 |
| Acrocephalus schoenobaenus | Denmark | -0.003 | 6.93 | 62.10 | 11.9 | 56.16 | 10.38 | 0.595 | -0.357 | 0 |
| Acrocephalus scirpaceus | Denmark | -0.009 | 6.59 | 44.60 | 11.8 | 56.16 | 10.38 | 0.659 | -0.330 | 0 |
| Aegitahlos caudatus | France | 0.000 | 5.27 | 0.00 | 8.8 | 46.71 | 1.72 | 0.489 | -0.385 | 0 |
| Aegithalos caudatus | Spain | -0.010 | 10.84 | 0.00 | 8.8 | 39.90 | -2.99 | 0.500 | -0.385 | 0 |
| Aegithalos caudatus | Hungary | 0.155 | 6.36 | 0.00 | 8.8 | 47.16 | 19.51 | 0.489 | -0.385 | 0 |
| Alauda arvensis | Denmark | -0.008 | 31.42 | 13.02 | 36.4 | 56.16 | 10.38 | 0.581 | -0.033 | 1 |
| Alauda arvensis | Poland | 0.017 | 45.44 | 13.02 | 36.4 | 51.92 | 19.13 | 0.533 | -0.033 | 1 |
| Alcedo atthis | France | -0.042 | 8.54 | 0.00 | 32.4 | 46.71 | 1.72 | 0.429 | -0.099 | 0 |
| Alectoris rufa | Spain | 0.004 | 37.08 | 0.00 | 477.5 | 39.90 | -2.99 | 0.732 | 0.303 | 1 |
| Anas platyrhynchos | Denmark | 0.048 | 28.76 | 8.13 | 1119.0 | 56.16 | 10.38 | 0.579 | 0.720 | 0 |
| Anas platyrhynchos | France | 0.012 | 4.81 | 8.13 | 1119.0 | 46.71 | 1.72 | 0.533 | 0.720 | 0 |
| Anas platyrhynchos | Norway | 0.123 | 11.11 | 8.13 | 1119.0 | 64.56 | 18.01 | 0.709 | 0.720 | 0 |
| Anas platyrhynchos | Poland | 0.014 | 88.00 | 8.13 | 1119.0 | 51.92 | 19.13 | 0.531 | 0.720 | 0 |
| Anser anser | Denmark | 0.107 | 180.00 | 12.28 | 3464.5 | 56.16 | 10.38 | 0.578 | 1.080 | 0 |
| Anthus pratensis | Denmark | -0.027 | 13.22 | 15.64 | 19.3 | 56.16 | 10.38 | 0.595 | -0.312 | 0 |
| Anthus pratensis | Finland | 0.002 | 28.31 | 15.64 | 19.3 | 64.95 | 26.07 | 0.730 | -0.312 | 0 |
| Anthus pratensis | Norway | 0.001 | 5.82 | 15.64 | 19.3 | 64.56 | 18.01 | 0.722 | -0.312 | 0 |
| Anthus pratensis | Poland | -0.046 | 32.00 | 15.64 | 19.3 | 51.92 | 19.13 | 0.619 | -0.312 | 0 |
| Anthus spinoletta | France | -0.037 | 7.50 | 21.27 | 21.5 | 46.71 | 1.72 | 0.549 | -0.237 | 0 |
| Anthus trivialis | Denmark | -0.013 | 10.90 | 47.07 | 23.4 | 56.16 | 10.38 | 0.587 | -0.210 | 0 |
| Apus apus | Denmark | 0.006 | 38.10 | 59.39 | 39.7 | 56.16 | 10.38 | 0.585 | -0.205 | 0 |
| Ardea cinerea | Denmark | 0.195 | 62.08 | 1.50 | 1433.0 | 56.16 | 10.38 | 0.330 | 0.903 | 0 |
| Ardea cinerea | France | 0.049 | 26.00 | 1.50 | 1433.0 | 46.71 | 1.72 | 0.289 | 0.903 | 0 |
| Ardea cinerea | Norway | 0.207 | 50.00 | 1.50 | 1433.0 | 64.56 | 18.01 | 0.596 | 0.903 | 0 |
| Athene noctua | Spain | -0.008 | 25.73 | 0.00 | 168.0 | 39.90 | -2.99 | 0.439 | 0.579 | 1 |
| Aythya fuligula | Denmark | 0.015 | 10.68 | 17.08 | 656.5 | 56.16 | 10.38 | 0.603 | 0.651 | 0 |
| Buteo buteo | Czech Rep. | 0.012 | 55.26 | 29.57 | 806.5 | 49.80 | 15.48 | 0.502 | 0.896 | 0 |
| Buteo buteo | Denmark | 0.033 | 60.01 | 29.57 | 806.5 | 56.16 | 10.38 | 0.591 | 0.896 | 0 |
| Carduelis cannabina | Czech Rep. | -0.011 | 14.93 | 4.11 | 19.0 | 49.80 | 15.48 | 0.547 | -0.187 | 0 |
| Carduelis cannabina | Denmark | -0.011 | 10.80 | 4.11 | 19.0 | 56.16 | 10.38 | 0.635 | -0.187 | 0 |
| Carduelis cannabina | Poland | -0.062 | 12.89 | 4.11 | 19.0 | 51.92 | 19.13 | 0.574 | -0.187 | 0 |
| Carduelis cannabina | Spain | -0.009 | 18.51 | 4.11 | 19.0 | 39.90 | -2.99 | 0.532 | -0.187 | 0 |
| Carduelis carduelis | Czech Rep. | -0.009 | 14.95 | 1.16 | 15.6 | 49.80 | 15.48 | 0.564 | -0.240 | 0 |
| Carduelis carduelis | Denmark | 0.169 | 10.77 | 1.16 | 15.6 | 56.16 | 10.38 | 0.667 | -0.240 | 0 |
| Carduelis carduelis | France | -0.002 | 11.00 | 1.16 | 15.6 | 46.71 | 1.72 | 0.524 | -0.240 | 0 |


| Carduelis carduelis | Spain | -0.022 | 11.52 | 1.16 | 15.6 | 39.90 | -2.99 | 0.490 | -0.240 | 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Carduelis carduelis | Hungary | 0.032 | 8.99 | 1.16 | 15.6 | 47.16 | 19.51 | 0.530 | -0.240 | 0 |
| Carduelis chloris | Czech Rep. | -0.021 | 12.98 | 1.34 | 27.7 | 49.80 | 15.48 | 0.507 | -0.057 | 0 |
| Carduelis chloris | Denmark | 0.036 | 6.53 | 1.34 | 27.7 | 56.16 | 10.38 | 0.581 | -0.057 | 0 |
| Carduelis chloris | Finland | 0.253 | 15.22 | 1.34 | 27.7 | 64.95 | 26.07 | 0.730 | -0.057 | 0 |
| Carduelis chloris | France | -0.020 | 7.65 | 1.34 | 27.7 | 46.71 | 1.72 | 0.502 | -0.057 | 0 |
| Carduelis chloris | Spain | 0.033 | 15.92 | 1.34 | 27.7 | 39.90 | -2.99 | 0.518 | -0.057 | 0 |
| Carduelis chloris | Hungary | 0.050 | 18.43 | 1.34 | 27.7 | 47.16 | 19.51 | 0.501 | -0.057 | 0 |
| Carduelis chloris | Norway | 0.022 | 5.88 | 1.34 | 27.7 | 64.56 | 18.01 | 0.720 | -0.057 | 0 |
| Carduelis chloris | Poland | 0.009 | 15.11 | 1.34 | 27.7 | 51.92 | 19.13 | 0.531 | -0.057 | 0 |
| Carduelis flammea | Denmark | -0.001 | 4.50 | 9.46 | 13.1 | 56.16 | 10.38 | 0.622 | -0.263 | 0 |
| Carduelis flammea | Norway | -0.027 | 12.00 | 9.46 | 13.1 | 64.56 | 18.01 | 0.653 | -0.263 | 0 |
| Carduelis spinus | Finland | 0.015 | 10.11 | 6.83 | 13.8 | 64.95 | 26.07 | 0.740 | -0.272 | 0 |
| Carpodacus erythrinus | Finland | -0.014 | 9.44 | 0.00 | 13.8 | 64.95 | 26.07 | 0.804 | -- | 0 |
| Certhia brachydactyla | France | 0.031 | 7.12 | 0.00 | 9.2 | 46.71 | 1.72 | 0.635 | -0.338 | 0 |
| Certhia brachydactyla | Spain | 0.024 | 11.15 | 0.00 | 9.2 | 39.90 | -2.99 | 0.603 | -0.338 | 0 |
| Certhia familiaris | Denmark | 0.015 | 4.47 | 0.00 | 9.2 | 56.16 | 10.38 | 0.582 | -0.297 | 0 |
| Cettia cetti | Spain | 0.000 | 26.84 | 2.11 | 14.1 | 39.90 | -2.99 | 0.564 | -- | 0 |
| Cisticola juncidis | Spain | -0.011 | 31.89 | 0.00 | 8.5 | 39.90 | -2.99 | 0.337 | -- | 0 |
| Clamator glandarius | Spain | 0.185 | 55.60 | 4.58 | 153.5 | 39.90 | -2.99 | 0.344 | 0.205 | 0 |
| Coccothraustes coccothraustes | Czech Rep. | -0.020 | 22.17 | 5.07 | 54.7 | 49.80 | 15.48 | 0.591 | 0.222 | 0 |
| Coccothraustes coccothraustes | France | 0.038 | 5.10 | 5.07 | 54.7 | 46.71 | 1.72 | 0.551 | 0.222 | 0 |
| Coccothraustes coccothraustes | Hungary | 0.038 | 24.00 | 5.07 | 54.7 | 47.16 | 19.51 | 0.556 | 0.222 | 0 |
| Columba livia | France | 0.024 | 8.00 | 0.00 | 261.0 | 46.71 | 1.72 | 0.403 | 0.303 | 1 |
| Columba livia | Spain | 0.007 | 30.87 | 0.00 | 261.0 | 39.90 | -2.99 | 0.291 | 0.303 | 1 |
| Columba livia | Hungary | 0.350 | 6.00 | 0.00 | 261.0 | 47.16 | 19.51 | 0.410 | 0.303 | 1 |
| Columba oenas | France | -0.012 | 17.75 | 3.45 | 494.5 | 46.71 | 1.72 | 0.546 | 0.333 | 1 |
| Columba palumbus | Czech Rep. | 0.028 | 27.26 | 2.03 | 494.5 | 49.80 | 15.48 | 0.521 | 0.365 | 1 |
| Columba palumbus | Denmark | 0.018 | 28.17 | 2.03 | 494.5 | 56.16 | 10.38 | 0.606 | 0.365 | 1 |
| Columba palumbus | Finland | 0.024 | 30.00 | 2.03 | 494.5 | 64.95 | 26.07 | 0.812 | 0.365 | 1 |
| Columba palumbus | France | 0.052 | 14.43 | 2.03 | 494.5 | 46.71 | 1.72 | 0.485 | 0.365 | 1 |
| Columba palumbus | Spain | 0.011 | 45.40 | 2.03 | 494.5 | 39.90 | -2.99 | 0.461 | 0.365 | 1 |
| Columba palumbus | Hungary | 0.008 | 21.47 | 2.03 | 494.5 | 47.16 | 19.51 | 0.490 | 0.365 | 1 |
| Columba palumbus | Norway | 0.016 | 7.00 | 2.03 | 494.5 | 64.56 | 18.01 | 0.796 | 0.365 | 1 |
| Columba palumbus | Poland | 0.018 | 60.41 | 2.03 | 494.5 | 51.92 | 19.13 | 0.548 | 0.365 | 1 |
| Corvus corax | Denmark | 0.293 | 78.06 | 0.00 | 1200.6 | 56.16 | 10.38 | 0.448 | 1.189 | 0 |
| Corvus cornix | Czech Rep. | -0.019 | 13.89 | 5.71 | 544.5 | 49.80 | 15.48 | 0.528 | 0.944 | 0 |
| Corvus cornix | Denmark | 0.011 | 41.15 | 5.71 | 544.5 | 56.16 | 10.38 | 0.579 | 0.944 | 0 |
| Corvus cornix | Finland | -0.013 | 31.69 | 5.71 | 544.5 | 64.95 | 26.07 | 0.717 | 0.944 | 0 |
| Corvus cornix | Hungary | 0.113 | 24.09 | 5.71 | 544.5 | 47.16 | 19.51 | 0.536 | 0.944 | 0 |
| Corvus cornix | Norway | 0.001 | 17.30 | 5.71 | 544.5 | 64.56 | 18.01 | 0.708 | 0.944 | 0 |
| Corvus corone | France | 0.001 | 20.41 | 5.71 | 544.5 | 46.71 | 1.72 | 0.537 | 0.944 | 0 |
| Corvus frugilegus | Denmark | 0.031 | 46.53 | 2.32 | 453.5 | 56.16 | 10.38 | 0.645 | 0.904 | 1 |
| Corvus monedula | Denmark | 0.000 | 32.11 | 0.29 | 249.0 | 56.16 | 10.38 | 0.635 | 0.660 | 1 |
| Corvus monedula | Finland | 0.168 | 60.00 | 0.29 | 249.0 | 64.95 | 26.07 | 0.872 | 0.660 | 1 |
| Corvus monedula | Spain | -0.030 | 46.86 | 0.29 | 249.0 | 39.90 | -2.99 | 0.573 | 0.660 | 1 |
| Corvus monedula | Poland | 0.019 | 18.23 | 0.29 | 249.0 | 51.92 | 19.13 | 0.576 | 0.660 | 1 |
| Cuculus canorus | Denmark | -0.010 | 21.72 | 49.38 | 120.5 | 56.16 | 10.38 | 0.562 | 0.164 | 0 |
| Cyanopica cyanus | Spain | 0.060 | 56.86 | 0.00 | 71.0 | 39.90 | -2.99 | 0.579 | 0.314 | 0 |
| Delichon urbica | Spain | 0.026 | 35.51 | 44.25 | 19.6 | 39.90 | -2.99 | 0.378 | -0.262 | 0 |
| Dendrocopos major | Czech Rep. | 0.018 | 18.56 | 0.00 | 89.7 | 49.80 | 15.48 | 0.498 | 0.405 | 0 |
| Dendrocopos major | Denmark | 0.005 | 14.20 | 0.00 | 89.7 | 56.16 | 10.38 | 0.581 | 0.405 | 0 |
| Dendrocopos major | France | 0.033 | 14.00 | 0.00 | 89.7 | 46.71 | 1.72 | 0.460 | 0.405 | 0 |
| Dendrocopos major | Spain | 0.069 | 58.14 | 0.00 | 89.7 | 39.90 | -2.99 | 0.411 | 0.405 | 0 |
| Dendrocopos major | Hungary | -0.014 | 32.98 | 0.00 | 89.7 | 47.16 | 19.51 | 0.466 | 0.405 | 0 |
| Dendrocopos major | Norway | 0.097 | 9.85 | 0.00 | 89.7 | 64.56 | 18.01 | 0.745 | 0.405 | 0 |
| Dendrocopos syriacus | Hungary | 0.023 | 13.99 | 0.00 | 76.8 | 47.16 | 19.51 | 0.726 | -- | 0 |
| Dryocopus martius | France | 0.069 | 38.71 | 0.00 | 273.0 | 46.71 | 1.72 | 0.517 | 0.870 | 0 |


| Dryocopus martius | Poland | 0.025 | 50.16 | 0.00 | 273.0 | 51.92 | 19.13 | 0.542 | 0.870 | 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Egretta garzetta | France | 0.041 | 24.50 | 7.53 | 532.5 | 46.71 | 1.72 | 0.400 | -- | 0 |
| Emberiza cirlus | Spain | -0.008 | 15.00 | 0.48 | 23.8 | 39.90 | -2.99 | 0.583 | -0.135 | 1 |
| Emberiza citrinella | Czech Rep. | -0.014 | 15.54 | 4.72 | 26.8 | 49.80 | 15.48 | 0.611 | -0.106 | 1 |
| Emberiza citrinella | Denmark | -0.016 | 9.79 | 4.72 | 26.8 | 56.16 | 10.38 | 0.600 | -0.106 | 1 |
| Emberiza citrinella | Finland | -0.002 | 10.43 | 4.72 | 26.8 | 64.95 | 26.07 | 0.743 | -0.106 | 1 |
| Emberiza citrinella | Norway | -0.023 | 8.20 | 4.72 | 26.8 | 64.56 | 18.01 | 0.734 | -0.106 | 1 |
| Emberiza citrinella | Poland | -0.016 | 3.08 | 4.72 | 26.8 | 51.92 | 19.13 | 0.597 | -0.106 | 1 |
| Emberiza schoeniclus | Czech Rep. | 0.001 | 28.07 | 10.52 | 18.8 | 49.80 | 15.48 | 0.510 | -0.167 | 0 |
| Emberiza schoeniclus | Denmark | -0.008 | 9.17 | 10.52 | 18.8 | 56.16 | 10.38 | 0.575 | -0.167 | 0 |
| Emberiza schoeniclus | Norway | -0.022 | 8.00 | 10.52 | 18.8 | 64.56 | 18.01 | 0.705 | -0.167 | 0 |
| Emberiza schoeniclus | Poland | 0.026 | 36.51 | 10.52 | 18.8 | 51.92 | 19.13 | 0.527 | -0.167 | 0 |
| Erithacus rubecula | Czech Rep. | -0.001 | 16.71 | 5.00 | 16.4 | 49.80 | 15.48 | 0.510 | -0.196 | 0 |
| Erithacus rubecula | Denmark | 0.006 | 8.91 | 5.00 | 16.4 | 56.16 | 10.38 | 0.585 | -0.196 | 0 |
| Erithacus rubecula | France | 0.023 | 5.19 | 5.00 | 16.4 | 46.71 | 1.72 | 0.503 | -0.196 | 0 |
| Erithacus rubecula | Norway | -0.001 | 5.47 | 5.00 | 16.4 | 64.56 | 18.01 | 0.727 | -0.196 | 0 |
| Erithacus rubecula | Spain | 0.015 | 2.24 | 5.00 | 16.4 | 39.90 | -2.99 | 0.519 | -0.196 | 0 |
| Falco tinnunculus | Czech Rep. | -0.003 | 50.58 | 5.60 | 174.5 | 49.80 | 15.48 | 0.228 | 0.570 | 1 |
| Falco tinnunculus | Denmark | 0.019 | 28.35 | 5.60 | 174.5 | 56.16 | 10.38 | 0.294 | 0.570 | 1 |
| Falco tinnunculus | France | -0.040 | 6.32 | 5.60 | 174.5 | 46.71 | 1.72 | 0.290 | 0.570 | 1 |
| Falco tinnunculus | Spain | -0.012 | 115.49 | 5.60 | 174.5 | 39.90 | -2.99 | 0.350 | 0.570 | 1 |
| Falco tinnunculus | Hungary | -0.007 | 25.00 | 5.60 | 174.5 | 47.16 | 19.51 | 0.284 | 0.570 | 1 |
| Ficedula hypoleuca | Denmark | -0.034 | 5.39 | 43.00 | 14.4 | 56.16 | 10.38 | 0.583 | -0.364 | 0 |
| Ficedula hypoleuca | Finland | 0.013 | 6.59 | 43.00 | 14.4 | 64.95 | 26.07 | 0.726 | -0.364 | 0 |
| Ficedula hypoleuca | Norway | -0.019 | 7.07 | 43.00 | 14.4 | 64.56 | 18.01 | 0.718 | -0.364 | 0 |
| Fringilla coelebs | Czech Rep. | -0.010 | 13.62 | 5.54 | 24.2 | 49.80 | 15.48 | 0.499 | -0.126 | 0 |
| Fringilla coelebs | Denmark | 0.011 | 9.39 | 5.54 | 24.2 | 56.16 | 10.38 | 0.571 | -0.126 | 0 |
| Fringilla coelebs | Finland | -0.002 | 8.50 | 5.54 | 24.2 | 64.95 | 26.07 | 0.710 | -0.126 | 0 |
| Fringilla coelebs | France | -0.003 | 6.55 | 5.54 | 24.2 | 46.71 | 1.72 | 0.500 | -0.126 | 0 |
| Fringilla coelebs | Spain | 0.047 | 17.26 | 5.54 | 24.2 | 39.90 | -2.99 | 0.515 | -0.126 | 0 |
| Fringilla coelebs | Norway | 0.008 | 7.30 | 5.54 | 24.2 | 64.56 | 18.01 | 0.701 | -0.126 | 0 |
| Fringilla coelebs | Poland | 0.002 | 8.76 | 5.54 | 24.2 | 51.92 | 19.13 | 0.522 | -0.126 | 0 |
| Fulica atra | Denmark | 0.010 | 19.87 | 3.32 | 732.5 | 56.16 | 10.38 | 0.158 | 0.484 | 0 |
| Galerida cristata | Spain | -0.013 | 36.40 | 0.00 | 44.7 | 39.90 | -2.99 | 0.417 | 0.033 | 1 |
| Gallinago gallinago | Denmark | -0.028 | 25.83 | 7.05 | 106.5 | 56.16 | 10.38 | 0.279 | 0.117 | 0 |
| Gallinago gallinago | Finland | -0.009 | 54.15 | 7.05 | 106.5 | 64.95 | 26.07 | 0.542 | 0.117 | 0 |
| Gallinula chloropus | Denmark | -0.015 | 20.00 | 0.00 | 348.5 | 56.16 | 10.38 | 0.400 | 0.296 | 0 |
| Gallinula chloropus | France | 0.018 | 10.32 | 0.00 | 348.5 | 46.71 | 1.72 | 0.263 | 0.296 | 0 |
| Garrulus glandarius | Czech Rep. | 0.041 | 39.15 | 0.00 | 161.7 | 49.80 | 15.48 | 0.484 | 0.605 | 0 |
| Garrulus glandarius | Denmark | 0.001 | 20.95 | 0.00 | 161.7 | 56.16 | 10.38 | 0.564 | 0.605 | 0 |
| Garrulus glandarius | France | 0.030 | 11.16 | 0.00 | 161.7 | 46.71 | 1.72 | 0.446 | 0.605 | 0 |
| Garrulus glandarius | Hungary | 0.011 | 27.87 | 0.00 | 161.7 | 47.16 | 19.51 | 0.452 | 0.605 | 0 |
| Grus grus | Poland | 0.051 | 100.00 | 39.10 | 4541.5 | 51.92 | 19.13 | 0.572 | 1.246 | 1 |
| Haematopus ostralegus | Denmark | -0.009 | 40.01 | 21.39 | 531.0 | 56.16 | 10.38 | 0.587 | 0.583 | 0 |
| Haematopus ostralegus | Norway | -0.010 | 19.00 | 21.39 | 531.0 | 64.56 | 18.01 | 0.714 | 0.583 | 0 |
| Hippolais icterina | Denmark | -0.034 | 7.76 | 71.34 | 13.3 | 56.16 | 10.38 | 0.601 | -0.293 | 0 |
| Hippolais polyglotta | Spain | 0.028 | 16.57 | 30.63 | 11.5 | 39.90 | -2.99 | 0.619 | -- | 0 |
| Hirundo rustica | Czech Rep. | -0.007 | 15.63 | 42.34 | 19.1 | 49.80 | 15.48 | 0.482 | -0.269 | 1 |
| Hirundo rustica | Denmark | -0.008 | 10.15 | 42.34 | 19.1 | 56.16 | 10.38 | 0.561 | -0.269 | 1 |
| Hirundo rustica | Finland | -0.013 | 11.18 | 42.34 | 19.1 | 64.95 | 26.07 | 0.712 | -0.269 | 1 |
| Hirundo rustica | Poland | 0.007 | 10.77 | 42.34 | 19.1 | 51.92 | 19.13 | 0.507 | -0.269 | 1 |
| Lanius collurio | Czech Rep. | 0.046 | 21.31 | 64.73 | 30.7 | 49.80 | 15.48 | 0.554 | 0.008 | 1 |
| Lanius collurio | Denmark | -0.011 | 17.39 | 64.73 | 30.7 | 56.16 | 10.38 | 0.639 | 0.008 | 1 |
| Lanius collurio | Hungary | 0.017 | 11.91 | 64.73 | 30.7 | 47.16 | 19.51 | 0.559 | 0.008 | 1 |
| Lanius collurio | Poland | -0.003 | 35.01 | 64.73 | 30.7 | 51.92 | 19.13 | 0.580 | 0.008 | 1 |
| Lanius excubitor | Poland | 0.072 | 22.36 | 4.87 | 66.9 | 51.92 | 19.13 | 0.555 | 0.199 | 1 |
| Lanius excubitor | Spain | -0.044 | 31.05 | 4.87 | 66.9 | 39.90 | -2.99 | 0.669 | 0.199 | 1 |
| Lanius senator | Spain | -0.012 | 36.12 | 28.45 | 36.0 | 39.90 | -2.99 | 0.575 | -- | 1 |


| Larus argentatus | Denmark | 0.019 | 50.09 | 22.14 | 895.0 | 56.16 | 10.38 | 0.673 | 0.774 | 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Larus argentatus | France | 0.050 | 15.10 | 22.14 | 895.0 | 46.71 | 1.72 | 0.952 | 0.774 | 0 |
| Larus canus | Denmark | -0.009 | 59.94 | 13.76 | 386.5 | 56.16 | 10.38 | 0.671 | 0.602 | 1 |
| Larus canus | Norway | -0.024 | 14.00 | 13.76 | 386.5 | 64.56 | 18.01 | 0.729 | 0.602 | 1 |
| Larus fuscus | France | 0.170 | 22.00 | 34.35 | 817.5 | 46.71 | 1.72 | 0.646 | 0.726 | 0 |
| Larus marinus | Denmark | 0.220 | 57.52 | 7.28 | 1599.5 | 56.16 | 10.38 | 0.528 | 0.837 | 0 |
| Larus ridibundus | Denmark | -0.033 | 37.75 | 23.50 | 280.5 | 56.16 | 10.38 | 0.624 | 0.453 | 0 |
| Locustella naevia | Czech Rep. | -0.010 | 7.22 | 29.75 | 12.7 | 49.80 | 15.48 | 0.597 | -0.222 | 1 |
| Loxia curvirostra | Denmark | -0.040 | 4.74 | 0.00 | 40.6 | 56.16 | 10.38 | 0.571 | 0.173 | 0 |
| Luscinia luscinia | Denmark | -0.016 | 15.89 | 63.10 | 25.0 | 56.16 | 10.38 | 0.670 | -0.125 | 0 |
| Luscinia megarhynchos | Czech Rep. | 0.099 | 15.52 | 32.20 | 20.2 | 49.80 | 15.48 | 0.722 | -0.179 | 0 |
| Luscinia megarhynchos | Spain | 0.018 | 25.87 | 32.20 | 20.2 | 39.90 | -2.99 | 0.560 | -0.179 | 0 |
| Luscinia megarhynchos | Hungary | 0.023 | 10.53 | 32.20 | 20.2 | 47.16 | 19.51 | 0.659 | -0.179 | 0 |
| Luscinia megarhynchos | Poland | -0.011 | 46.01 | 32.20 | 20.2 | 51.92 | 19.13 | 0.794 | -0.179 | 0 |
| Merops apiaster | Spain | 0.008 | 94.95 | 19.86 | 55.1 | 39.90 | -2.99 | 0.335 | -0.098 | 1 |
| Miliaria calandra | Denmark | -0.004 | 10.29 | 6.40 | 47.7 | 56.16 | 10.38 | 0.782 | 0.074 | 1 |
| Miliaria calandra | Spain | -0.008 | 24.18 | 6.40 | 47.7 | 39.90 | -2.99 | 0.576 | 0.074 | 1 |
| Miliaria calandra | Poland | 0.064 | 31.64 | 6.40 | 47.7 | 51.92 | 19.13 | 0.671 | 0.074 | 1 |
| Motacilla alba | Czech Rep. | -0.014 | 18.03 | 18.13 | 20.8 | 49.80 | 15.48 | 0.480 | -0.250 | 0 |
| Motacilla alba | Denmark | 0.049 | 11.62 | 18.13 | 20.8 | 56.16 | 10.38 | 0.556 | -0.250 | 0 |
| Motacilla alba | Finland | 0.000 | 7.57 | 18.13 | 20.8 | 64.95 | 26.07 | 0.699 | -0.250 | 0 |
| Motacilla alba | France | 0.002 | 6.33 | 18.13 | 20.8 | 46.71 | 1.72 | 0.445 | -0.250 | 0 |
| Motacilla alba | Spain | -0.019 | 24.36 | 18.13 | 20.8 | 39.90 | -2.99 | 0.421 | -0.250 | 0 |
| Motacilla alba | Norway | 0.026 | 7.15 | 18.13 | 20.8 | 64.56 | 18.01 | 0.690 | -0.250 | 0 |
| Motacilla alba | Poland | 0.005 | 25.56 | 18.13 | 20.8 | 51.92 | 19.13 | 0.505 | -0.250 | 0 |
| Motacilla cinerea | Czech Rep. | -0.017 | 8.00 | 24.68 | 17.4 | 49.80 | 15.48 | 0.507 | -0.279 | 0 |
| Motacilla cinerea | Denmark | -0.027 | 6.87 | 24.68 | 17.4 | 56.16 | 10.38 | 0.562 | -0.279 | 0 |
| Motacilla cinerea | France | -0.015 | 9.07 | 24.68 | 17.4 | 46.71 | 1.72 | 0.503 | -0.279 | 0 |
| Motacilla flava | Denmark | -0.027 | 11.20 | 40.99 | 17.5 | 56.16 | 10.38 | 0.562 | -0.371 | 1 |
| Motacilla flava | Finland | -0.036 | 11.84 | 40.99 | 17.5 | 64.95 | 26.07 | 0.709 | -0.371 | 1 |
| Motacilla flava | Poland | -0.030 | 5.39 | 40.99 | 17.5 | 51.92 | 19.13 | 0.509 | -0.371 | 1 |
| Muscicapa striata | Czech Rep. | 0.057 | 15.56 | 64.40 | 15.5 | 49.80 | 15.48 | 0.516 | -0.292 | 0 |
| Muscicapa striata | Denmark | 0.010 | 10.08 | 64.40 | 15.5 | 56.16 | 10.38 | 0.584 | -0.292 | 0 |
| Muscicapa striata | Finland | -0.004 | 9.06 | 64.40 | 15.5 | 64.95 | 26.07 | 0.729 | -0.292 | 0 |
| Muscicapa striata | Spain | -0.025 | 21.98 | 64.40 | 15.5 | 39.90 | -2.99 | 0.551 | -0.292 | 0 |
| Numenius arquata | Finland | -0.008 | 42.07 | 44.67 | 725.0 | 64.95 | 26.07 | 0.744 | 0.595 | 0 |
| Oriolus oriolus | Spain | 0.056 | 30.34 | 44.77 | 68.5 | 39.90 | -2.99 | 0.485 | 0.117 | 0 |
| Oriolus oriolus | Hungary | 0.003 | 24.12 | 44.77 | 68.5 | 47.16 | 19.51 | 0.536 | 0.117 | 0 |
| Parus ater | Czech Rep. | 0.017 | 12.08 | 0.00 | 9.3 | 49.80 | 15.48 | 0.520 | -0.264 | 0 |
| Parus ater | Denmark | -0.009 | 5.41 | 0.00 | 9.3 | 56.16 | 10.38 | 0.605 | -0.264 | 0 |
| Parus ater | France | -0.049 | 4.00 | 0.00 | 9.3 | 46.71 | 1.72 | 0.483 | -0.264 | 0 |
| Parus ater | Spain | 0.019 | 18.49 | 0.00 | 9.3 | 39.90 | -2.99 | 0.456 | -0.264 | 0 |
| Parus caeruleus | Czech Rep. | 0.002 | 10.08 | 0.00 | 11.8 | 49.80 | 15.48 | 0.540 | -0.200 | 0 |
| Parus caeruleus | Denmark | 0.007 | 6.61 | 0.00 | 11.8 | 56.16 | 10.38 | 0.622 | -0.200 | 0 |
| Parus caeruleus | Finland | 0.258 | 8.00 | 0.00 | 11.8 | 64.95 | 26.07 | 0.825 | -0.200 | 0 |
| Parus caeruleus | France | 0.033 | 5.17 | 0.00 | 11.8 | 46.71 | 1.72 | 0.537 | -0.200 | 0 |
| Parus caeruleus | Spain | 0.027 | 9.93 | 0.00 | 11.8 | 39.90 | -2.99 | 0.570 | -0.200 | 0 |
| Parus caeruleus | Hungary | 0.025 | 5.94 | 0.00 | 11.8 | 47.16 | 19.51 | 0.535 | -0.200 | 0 |
| Parus caeruleus | Norway | 0.012 | 6.09 | 0.00 | 11.8 | 64.56 | 18.01 | 0.808 | -0.200 | 0 |
| Parus cristatus | Denmark | -0.013 | 7.20 | 0.00 | 11.2 | 56.16 | 10.38 | 0.636 | -0.189 | 0 |
| Parus cristatus | France | 0.005 | 4.81 | 0.00 | 11.2 | 46.71 | 1.72 | 0.603 | -0.189 | 0 |
| Parus major | Czech Rep. | -0.002 | 10.47 | 0.00 | 18.5 | 49.80 | 15.48 | 0.370 | -0.073 | 0 |
| Parus major | Denmark | -0.009 | 5.46 | 0.00 | 18.5 | 56.16 | 10.38 | 0.479 | -0.073 | 0 |
| Parus major | Finland | 0.021 | 8.61 | 0.00 | 18.5 | 64.95 | 26.07 | 0.649 | -0.073 | 0 |
| Parus major | France | 0.017 | 4.60 | 0.00 | 18.5 | 46.71 | 1.72 | 0.307 | -0.073 | 0 |
| Parus major | Spain | 0.013 | 11.92 | 0.00 | 18.5 | 39.90 | -2.99 | 0.000 | -0.073 | 0 |
| Parus major | Hungary | 0.027 | 7.65 | 0.00 | 18.5 | 47.16 | 19.51 | 0.317 | -0.073 | 0 |
| Parus major | Norway | 0.003 | 5.74 | 0.00 | 18.5 | 64.56 | 18.01 | 0.639 | -0.073 | 0 |


| Parus major | Poland | -0.009 | 16.49 | 0.00 | 18.5 | 51.92 | 19.13 | 0.408 | -0.073 | 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Parus montanus | Norway | -0.004 | 5.61 | 0.00 | 11.7 | 64.56 | 18.01 | 0.716 | -0.102 | 0 |
| Parus palustris | Czech Rep. | -0.022 | 10.60 | 0.00 | 11.9 | 49.80 | 15.48 | 0.542 | -0.151 | 0 |
| Parus palustris | Denmark | -0.013 | 4.03 | 0.00 | 11.9 | 56.16 | 10.38 | 0.638 | -0.151 | 0 |
| Parus palustris | France | -0.008 | 6.60 | 0.00 | 11.9 | 46.71 | 1.72 | 0.503 | -0.151 | 0 |
| Passer domesticus | Czech Rep. | -0.026 | 15.28 | 0.00 | 30.4 | 49.80 | 15.48 | 0.435 | -0.041 | 0 |
| Passer domesticus | Denmark | -0.013 | 4.72 | 0.00 | 30.4 | 56.16 | 10.38 | 0.521 | -0.041 | 0 |
| Passer domesticus | Finland | -0.038 | 16.67 | 0.00 | 30.4 | 64.95 | 26.07 | 0.671 | -0.041 | 0 |
| Passer domesticus | France | 0.004 | 4.83 | 0.00 | 30.4 | 46.71 | 1.72 | 0.392 | -0.041 | 0 |
| Passer domesticus | Spain | -0.004 | 19.76 | 0.00 | 30.4 | 39.90 | -2.99 | 0.277 | -0.041 | 0 |
| Passer domesticus | Hungary | -0.004 | 7.34 | 0.00 | 30.4 | 47.16 | 19.51 | 0.398 | -0.041 | 0 |
| Passer domesticus | Norway | -0.008 | 4.24 | 0.00 | 30.4 | 64.56 | 18.01 | 0.662 | -0.041 | 0 |
| Passer domesticus | Poland | -0.027 | 11.05 | 0.00 | 30.4 | 51.92 | 19.13 | 0.463 | -0.041 | 0 |
| Passer montanus | Czech Rep. | -0.009 | 16.67 | 3.62 | 21.7 | 49.80 | 15.48 | 0.378 | -0.123 | 1 |
| Passer montanus | Denmark | 0.036 | 6.35 | 3.62 | 21.7 | 56.16 | 10.38 | 0.487 | -0.123 | 1 |
| Passer montanus | Spain | -0.027 | 13.79 | 3.62 | 21.7 | 39.90 | -2.99 | 0.041 | -0.123 | 1 |
| Passer montanus | Hungary | 0.073 | 8.08 | 3.62 | 21.7 | 47.16 | 19.51 | 0.325 | -0.123 | 1 |
| Passer montanus | Poland | -0.051 | 14.45 | 3.62 | 21.7 | 51.92 | 19.13 | 0.415 | -0.123 | 1 |
| Perdix perdix | Denmark | -0.012 | 24.92 | 0.00 | 382.0 | 56.16 | 10.38 | 0.655 | 0.259 | 1 |
| Phoenicurus ochruros | Czech Rep. | 0.009 | 15.89 | 15.83 | 16.0 | 49.80 | 15.48 | 0.612 | -0.254 | 0 |
| Phoenicurus ochruros | France | 0.006 | 4.12 | 15.83 | 16.0 | 46.71 | 1.72 | 0.567 | -0.254 | 0 |
| Phoenicurus ochruros | Hungary | 0.061 | 15.10 | 15.83 | 16.0 | 47.16 | 19.51 | 0.573 | -0.254 | 0 |
| Phoenicurus ochruros | Poland | -0.011 | 22.40 | 15.83 | 16.0 | 51.92 | 19.13 | 0.648 | -0.254 | 0 |
| Phoenicurus phoenicurus | Czech Rep. | 0.061 | 19.10 | 33.93 | 15.9 | 49.80 | 15.48 | 0.505 | -0.321 | 0 |
| Phoenicurus phoenicurus | Denmark | 0.019 | 12.05 | 33.93 | 15.9 | 56.16 | 10.38 | 0.579 | -0.321 | 0 |
| Phoenicurus phoenicurus | Finland | 0.020 | 20.02 | 33.93 | 15.9 | 64.95 | 26.07 | 0.725 | -0.321 | 0 |
| Phylloscopus collybita | Czech Rep. | 0.010 | 11.03 | 22.55 | 7.7 | 49.80 | 15.48 | 0.507 | -0.455 | 0 |
| Phylloscopus collybita | Denmark | 0.168 | 7.88 | 22.55 | 7.7 | 56.16 | 10.38 | 0.581 | -0.455 | 0 |
| Phylloscopus collybita | France | -0.018 | 5.25 | 22.55 | 7.7 | 46.71 | 1.72 | 0.500 | -0.455 | 0 |
| Phylloscopus trochilus | Denmark | -0.020 | 6.06 | 68.09 | 9.4 | 56.16 | 10.38 | 0.617 | -0.507 | 0 |
| Phylloscopus trochilus | Hungary | -0.034 | 2.00 | 68.09 | 9.4 | 47.16 | 19.51 | 0.725 | -0.507 | 0 |
| Phylloscopus trochilus | Norway | -0.011 | 5.05 | 68.09 | 9.4 | 64.56 | 18.01 | 0.725 | -0.507 | 0 |
| Pica pica | Denmark | 0.008 | 37.59 | 0.00 | 228.0 | 56.16 | 10.38 | 0.551 | 0.736 | 0 |
| Pica pica | Finland | 0.002 | 29.00 | 0.00 | 228.0 | 64.95 | 26.07 | 0.694 | 0.736 | 0 |
| Pica pica | France | -0.112 | 15.12 | 0.00 | 228.0 | 46.71 | 1.72 | 0.438 | 0.736 | 0 |
| Pica pica | Spain | -0.008 | 38.13 | 0.00 | 228.0 | 39.90 | -2.99 | 0.399 | 0.736 | 0 |
| Pica pica | Hungary | 0.016 | 21.46 | 0.00 | 228.0 | 47.16 | 19.51 | 0.443 | 0.736 | 0 |
| Pica pica | Norway | 0.000 | 13.98 | 0.00 | 228.0 | 64.56 | 18.01 | 0.685 | 0.736 | 0 |
| Pica pica | Poland | 0.022 | 26.66 | 0.00 | 228.0 | 51.92 | 19.13 | 0.499 | 0.736 | 0 |
| Picus viridis | France | 0.026 | 17.82 | 0.00 | 193.5 | 46.71 | 1.72 | 0.587 | 0.636 | 0 |
| Picus viridis | Spain | -0.006 | 50.24 | 0.00 | 193.5 | 39.90 | -2.99 | 0.657 | 0.636 | 0 |
| Pluvialis apricaria | Finland | -0.010 | 18.00 | 17.97 | 175.5 | 64.95 | 26.07 | 0.741 | 0.314 | 0 |
| Prunella modularis | Czech Rep. | -0.011 | 13.56 | 9.23 | 19.0 | 49.80 | 15.48 | 0.585 | -0.163 | 0 |
| Prunella modularis | Denmark | -0.018 | 7.43 | 9.23 | 19.0 | 56.16 | 10.38 | 0.594 | -0.163 | 0 |
| Prunella modularis | France | -0.006 | 4.61 | 9.23 | 19.0 | 46.71 | 1.72 | 0.606 | -0.163 | 0 |
| Pyrrhula pyrrhula | Denmark | 0.010 | 7.65 | 0.00 | 31.1 | 56.16 | 10.38 | 0.592 | -0.058 | 0 |
| Pyrrhula pyrrhula | Finland | 0.061 | 5.00 | 0.00 | 31.1 | 64.95 | 26.07 | 0.738 | -0.058 | 0 |
| Pyrrhula pyrrhula | Norway | -0.042 | 16.00 | 0.00 | 31.1 | 64.56 | 18.01 | 0.729 | -0.058 | 0 |
| Regulus regulus | Denmark | -0.008 | 5.41 | 0.00 | 5.8 | 56.16 | 10.38 | 0.578 | -0.446 | 0 |
| Regulus regulus | France | -0.003 | 3.24 | 0.00 | 5.8 | 46.71 | 1.72 | 0.484 | -0.446 | 0 |
| Regulus regulus | Norway | -0.007 | 5.11 | 0.00 | 5.8 | 64.56 | 18.01 | 0.718 | -0.446 | 0 |
| Riparia riparia | Denmark | -0.020 | 23.10 | 42.73 | 13.2 | 56.16 | 10.38 | 0.572 | -0.466 | 0 |
| Saxicola rubetra | Denmark | -0.040 | 15.43 | 34.84 | 16.6 | 56.16 | 10.38 | 0.602 | -0.222 | 1 |
| Saxicola rubetra | Finland | -0.018 | 23.09 | 34.84 | 16.6 | 64.95 | 26.07 | 0.751 | -0.222 | 1 |
| Saxicola torquata | Hungary | -0.006 | 15.98 | 3.98 | 14.9 | 47.16 | 19.51 | 0.280 | -- | 1 |
| Scolopax rusticola | Finland | 0.020 | 9.00 | 14.32 | 309.5 | 64.95 | 26.07 | 0.732 | 0.388 | 0 |
| Serinus serinus | Czech Rep. | -0.039 | 13.94 | 4.91 | 12.0 | 49.80 | 15.48 | 0.621 | -0.371 | 0 |
| Serinus serinus | France | -0.033 | 6.06 | 4.91 | 12.0 | 46.71 | 1.72 | 0.574 | -0.371 | 0 |


| Serinus serinus | Spain | -0.025 | 21.67 | 4.91 | 12.0 | 39.90 | -2.99 | 0.551 | -0.371 | 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Serinus serinus | Hungary | -0.059 | 14.14 | 4.91 | 12.0 | 47.16 | 19.51 | 0.581 | -0.371 | 0 |
| Serinus serinus | Poland | -0.044 | 24.04 | 4.91 | 12.0 | 51.92 | 19.13 | 0.658 | -0.371 | 0 |
| Sitta europaea | Czech Rep. | 0.014 | 13.39 | 0.00 | 23.9 | 49.80 | 15.48 | 0.492 | 0.017 | 0 |
| Sitta europaea | France | -0.038 | 4.92 | 0.00 | 23.9 | 46.71 | 1.72 | 0.450 | 0.017 | 0 |
| Sitta europaea | Norway | 0.025 | 5.41 | 0.00 | 23.9 | 64.56 | 18.01 | 0.788 | 0.017 | 0 |
| Streptopelia decaocto | Czech Rep. | 0.026 | 22.42 | 0.00 | 201.5 | 49.80 | 15.48 | 0.460 | 0.157 | 0 |
| Streptopelia decaocto | Denmark | 0.000 | 26.43 | 0.00 | 201.5 | 56.16 | 10.38 | 0.547 | 0.157 | 0 |
| Streptopelia decaocto | France | 0.077 | 11.67 | 0.00 | 201.5 | 46.71 | 1.72 | 0.418 | 0.157 | 0 |
| Streptopelia decaocto | Spain | 0.538 | 23.61 | 0.00 | 201.5 | 39.90 | -2.99 | 0.315 | 0.157 | 0 |
| Streptopelia decaocto | Hungary | 0.044 | 10.79 | 0.00 | 201.5 | 47.16 | 19.51 | 0.424 | 0.157 | 0 |
| Streptopelia decaocto | Poland | 0.005 | 13.31 | 0.00 | 201.5 | 51.92 | 19.13 | 0.488 | 0.157 | 0 |
| Streptopelia turtur | Czech Rep. | -0.022 | 27.99 | 25.90 | 136.5 | 49.80 | 15.48 | 0.550 | 0.102 | 1 |
| Streptopelia turtur | France | -0.006 | 7.07 | 25.90 | 136.5 | 46.71 | 1.72 | 0.509 | 0.102 | 1 |
| Streptopelia turtur | Spain | -0.004 | 97.53 | 25.90 | 136.5 | 39.90 | -2.99 | 0.440 | 0.102 | 1 |
| Sturnus unicolor | Spain | 0.018 | 42.70 | 0.00 | 90.6 | 39.90 | -2.99 | 0.728 | 0.262 | 1 |
| Sturnus vulgaris | Czech Rep. | 0.030 | 36.44 | 2.63 | 80.5 | 49.80 | 15.48 | 0.493 | 0.235 | 1 |
| Sturnus vulgaris | Denmark | -0.015 | 14.31 | 2.63 | 80.5 | 56.16 | 10.38 | 0.567 | 0.235 | 1 |
| Sturnus vulgaris | France | -0.005 | 9.75 | 2.63 | 80.5 | 46.71 | 1.72 | 0.477 | 0.235 | 1 |
| Sturnus vulgaris | Hungary | 0.042 | 13.46 | 2.63 | 80.5 | 47.16 | 19.51 | 0.477 | 0.235 | 1 |
| Sturnus vulgaris | Norway | -0.016 | 8.83 | 2.63 | 80.5 | 64.56 | 18.01 | 0.697 | 0.235 | 1 |
| Sturnus vulgaris | Poland | 0.049 | 31.28 | 2.63 | 80.5 | 51.92 | 19.13 | 0.517 | 0.235 | 1 |
| Sylvia atricapilla | Czech Rep. | 0.074 | 13.56 | 19.63 | 18.9 | 49.80 | 15.48 | 0.476 | -0.207 | 0 |
| Sylvia atricapilla | Denmark | 0.044 | 8.48 | 19.63 | 18.9 | 56.16 | 10.38 | 0.557 | -0.207 | 0 |
| Sylvia atricapilla | France | 0.013 | 4.83 | 19.63 | 18.9 | 46.71 | 1.72 | 0.438 | -0.207 | 0 |
| Sylvia atricapilla | Spain | 0.044 | 11.47 | 19.63 | 18.9 | 39.90 | -2.99 | 0.373 | -0.207 | 0 |
| Sylvia atricapilla | Hungary | 0.056 | 10.81 | 19.63 | 18.9 | 47.16 | 19.51 | 0.443 | -0.207 | 0 |
| Sylvia atricapilla | Norway | 0.119 | 3.16 | 19.63 | 18.9 | 64.56 | 18.01 | 0.704 | -0.207 | 0 |
| Sylvia atricapilla | Poland | 0.031 | 11.32 | 19.63 | 18.9 | 51.92 | 19.13 | 0.502 | -0.207 | 0 |
| Sylvia borin | Czech Rep. | -0.012 | 19.10 | 63.25 | 19.1 | 49.80 | 15.48 | 0.584 | -0.220 | 0 |
| Sylvia borin | Denmark | -0.014 | 6.78 | 63.25 | 19.1 | 56.16 | 10.38 | 0.598 | -0.220 | 0 |
| Sylvia cantillans | Spain | 0.028 | 5.22 | 20.82 | 8.1 | 39.90 | -2.99 | 0.668 | -- | 0 |
| Sylvia communis | Czech Rep. | 0.000 | 14.70 | 53.05 | 14.5 | 49.80 | 15.48 | 0.527 | -0.281 | 1 |
| Sylvia communis | Denmark | 0.004 | 7.54 | 53.05 | 14.5 | 56.16 | 10.38 | 0.598 | -0.281 | 1 |
| Sylvia communis | Spain | -0.021 | 22.94 | 53.05 | 14.5 | 39.90 | -2.99 | 0.570 | -0.281 | 1 |
| Sylvia curruca | Czech Rep. | 0.041 | 17.26 | 27.79 | 12.4 | 49.80 | 15.48 | 0.514 | -0.296 | 0 |
| Sylvia curruca | Denmark | -0.022 | 5.72 | 27.79 | 12.4 | 56.16 | 10.38 | 0.590 | -0.296 | 0 |
| Sylvia curruca | Hungary | -0.007 | 8.55 | 27.79 | 12.4 | 47.16 | 19.51 | 0.509 | -0.296 | 0 |
| Sylvia melanocephala | Spain | -0.002 | 23.22 | 3.61 | 13.5 | 39.90 | -2.99 | 0.660 | -0.269 | 0 |
| Tadorna tadorna | Denmark | 0.013 | 37.71 | 9.35 | 1152.0 | 56.16 | 10.38 | 0.638 | 0.684 | 0 |
| Tringa hypoleucos | France | -0.021 | 20.00 | 44.39 | 47.8 | 46.71 | 1.72 | 0.527 | -0.108 | 0 |
| Tringa totanus | Denmark | -0.030 | 29.71 | 35.28 | 112.0 | 56.16 | 10.38 | 0.576 | 0.149 | 0 |
| Troglodytes troglodytes | Czech Rep. | 0.011 | 10.55 | 1.34 | 8.9 | 49.80 | 15.48 | 0.497 | -0.312 | 0 |
| Troglodytes troglodytes | Denmark | 0.029 | 7.51 | 1.34 | 8.9 | 56.16 | 10.38 | 0.577 | -0.312 | 0 |
| Troglodytes troglodytes | France | 0.012 | 4.90 | 1.34 | 8.9 | 46.71 | 1.72 | 0.461 | -0.312 | 0 |
| Troglodytes troglodytes | Norway | -0.002 | 3.00 | 1.34 | 8.9 | 64.56 | 18.01 | 0.731 | -0.312 | 0 |
| Turdus iliacus | Finland | -0.002 | 14.05 | 10.77 | 62.9 | 64.95 | 26.07 | 0.741 | 0.133 | 0 |
| Turdus iliacus | Norway | 0.026 | 4.00 | 10.77 | 62.9 | 64.56 | 18.01 | 0.732 | 0.133 | 0 |
| Turdus merula | Czech Rep. | 0.016 | 21.51 | 3.98 | 95.9 | 49.80 | 15.48 | 0.438 | 0.284 | 0 |
| Turdus merula | Denmark | 0.015 | 12.22 | 3.98 | 95.9 | 56.16 | 10.38 | 0.524 | 0.284 | 0 |
| Turdus merula | France | 0.016 | 8.02 | 3.98 | 95.9 | 46.71 | 1.72 | 0.396 | 0.284 | 0 |
| Turdus merula | Spain | 0.007 | 21.98 | 3.98 | 95.9 | 39.90 | -2.99 | 0.284 | 0.284 | 0 |
| Turdus merula | Hungary | 0.012 | 9.08 | 3.98 | 95.9 | 47.16 | 19.51 | 0.402 | 0.284 | 0 |
| Turdus merula | Norway | 0.015 | 9.65 | 3.98 | 95.9 | 64.56 | 18.01 | 0.665 | 0.284 | 0 |
| Turdus merula | Poland | -0.018 | 22.59 | 3.98 | 95.9 | 51.92 | 19.13 | 0.466 | 0.284 | 0 |
| Turdus philomelos | Czech Rep. | -0.001 | 29.03 | 14.65 | 70.5 | 49.80 | 15.48 | 0.576 | 0.196 | 0 |
| Turdus philomelos | Denmark | -0.004 | 15.37 | 14.65 | 70.5 | 56.16 | 10.38 | 0.596 | 0.196 | 0 |
| Turdus philomelos | France | 0.034 | 8.10 | 14.65 | 70.5 | 46.71 | 1.72 | 0.595 | 0.196 | 0 |


| Turdus pilaris | Czech Rep. | -0.003 | 29.31 | 10.77 | 92.1 | 49.80 | 15.48 | 0.709 | 0.261 |  |
| :--- | :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Turdus pilaris | Denmark | -0.001 | 18.14 | 10.77 | 92.1 | 56.16 | 10.38 | 0.634 | 0.261 |  |
| Turdus pilaris | Finland | 0.037 | 20.95 | 10.77 | 92.1 | 64.95 | 26.07 | 0.735 | 0.261 | 1 |
| Turdus pilaris | Norway | -0.015 | 8.56 | 10.77 | 92.1 | 64.56 | 18.01 | 0.727 | 0.261 |  |
| Turdus pilaris | Poland | -0.038 | 5.50 | 10.77 | 92.1 | 51.92 | 19.13 | 0.677 | 0.261 |  |
| Turdus viscivorus | Czech Rep. | 0.024 | 49.90 | 4.36 | 117.8 | 49.80 | 15.48 | 0.526 | 0.313 |  |
| Turdus viscivorus | Denmark | 0.018 | 23.86 | 4.36 | 117.8 | 56.16 | 10.38 | 0.602 | 0.313 | 0 |
| Turdus viscivorus | France | -0.011 | 27.00 | 4.36 | 117.8 | 46.71 | 1.72 | 0.534 | 0.313 | 0 |
| Upupa epops | Spain | 0.001 | 40.43 | 9.59 | 67.1 | 39.90 | -2.99 | 0.339 | 0.102 | 0 |
| Vanellus vanellus | Denmark | -0.011 | 33.99 | 12.08 | 218.5 | 56.16 | 10.38 | 0.598 | 0.326 |  |

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