INTRODUCTION

Recent global assessments of environmental change highlight human-driven loss of biodiversity and the degradation of ecosystem integrity (Díaz et al., 2019; Secretariat of the Convention on Biological Diversity, 2020). Further, they point to a failure to achieve existing biodiversity targets and call for transformative change across sectors of human society as an emerging Post-2020 Global Biodiversity Framework takes shape (https://www.cbd.int/).

Yet at the same time, debate continues as to the nature and extent of biodiversity decline (e.g., Dornelas et al., 2019; Gonzalez et al., 2016; Leung et al., 2020; Vellend et al., 2017). That contrast in part reflects the inherent complexity of biodiversity and how temporal change can be measured. Different metrics capture different dimensions of biodiversity, are not necessarily correlated with each other, and differ in their suitability to measure biodiversity change, and some perform consistently better than others in that respect (Santini et al., 2017; Schipper et al., 2016). Recent attention has focused on the underlying pattern of change across species and on the extremes (Leung et al., 2020).

Although there is evidence for widespread declines from population surveys, assemblage surveys tend to suggest a more balanced picture of change (Dornelas et al., 2019; Vellend et al., 2013, 2017). To help understand those differences, Dornelas et al. (2019)
suggested that the selection of study populations may be inadvertently biased toward declining species and questioned whether such "ideal" datasets do, or would ever, exist. Buckland and Johnston (2017) argue for monitoring programs to have (1) representative sampling locations; (2) sufficient sample sizes; (3) sufficient detections of target species; (4) a representative sample of species; and (5) a temporal sampling scheme designed to aid valid inference. In reality, few individual schemes meet all of these criteria, and this issue becomes compounded when data from multiple schemes are integrated to assess population change. Combining datasets can mean that a more representative sample of species is included but may introduce other potential biases, for example, due to differences in temporal coverage or sampling size.

Population studies of birds are some of the most well-developed, reflecting the popularity of the taxa and the relative ease by which they are detected and identified (Gregory & van Strien, 2010). There is a rich tradition of bird surveys and atlas projects across the globe, often involving skilled amateur ornithologists with professionals in structured and well-designed monitoring projects (Harris et al., 2020; Keller et al., 2020). Taking advantage of extensive high-quality data, notably in the form of the North American Breeding Bird Survey, Rosenberg et al. (2019) modeled population change in 529 North American species (76% of breeding species) over 48 years. They integrated species' population trajectories and population size estimates into a hierarchical Bayesian model to produce a time series of population sizes across species, estimating a decline of 29% in breeding bird abundance since 1970 and indicating a staggering net loss of 2.9 billion birds. Using the same data, Jörger-Hickfang et al., 2020 estimated the proportional change per species and found a smaller average decline and suggested that biodiversity assessments should present a range, rather than a single measure of change.

Previous work using extensive data collected in Europe has also demonstrated substantial population declines in common and widespread birds, especially in those associated with agricultural systems (e.g., Donald et al., 2006; Gregory et al., 2005; Reif, 2013; Tucker & Heath, 1994) and in long-distance migrants (Sanderson et al., 2006, 2016). Inger et al. (2015) combined annual trend estimates with estimates of population size for 144 widespread European birds (32% of native breeding species) between 1980 and 2009 to demonstrate a significant decline in the total bird abundance and biomass. Most of this loss was attributed to the more common species, whereas less abundant species showed an overall increase in both abundance and biomass.

Recent work has also emphasized the importance of species abundance as a key component and metric of biodiversity change. This is both from a perspective reflecting the intrinsic value of species and their persistence and, more broadly, the fundamental role that species populations play in the functioning of ecological systems and in the provision of ecosystem services, or goods, upon which humanity relies (Gaston, 2010; Mace et al., 2012). In that respect, common species are likely to contribute disproportionately more than rare ones as even relatively small proportional declines in the abundance of common species will result in large absolute losses of individuals and biomass, which may disrupt ecosystem structure, function, and services (Gaston & Fuller, 2008). For that reason, ambitions to recover depleted populations and restore the abundance of species are increasingly prominent in national and international environmental frameworks, for example, within the draft Post-2020 Global Biodiversity Framework (United Nations Environment Programme, 2021).

In this study, we bring together two large avian datasets to explore abundance change in the European avifauna, taking the European Union (EU) as our geopolitical unit. Integrating these datasets allowed us to more than double the number of species included and to extend the temporal coverage by close to a decade compared with previous similar work. Our specific aim was to estimate change in the total population size of wild native breeding bird species in the EU and their average population growth rate, between 1980 and 2017, and to explore whether the previously observed patterns of change held true, given the extended taxonomic and temporal coverage.

We first test the hypothesis that the total avian abundance in the EU has not changed in the past 40 years, and whether the average population growth rate (log-transformed) is negative. We then test the degree to which these patterns are associated with the abundance class of the bird species, their habitat affinities, and aspects of their ecology. We predict that (1) the overall avian abundance in the EU has fallen and the average population growth rate is negative; (2) that this decline is most marked in abundant bird species; (3) that declines have been most pronounced in birds associated with particular habitats (e.g., agricultural landscapes); and (4) that aspects of ecology, such as migration strategy, are associated with trends. Across species and categories, we examine the overall numerical change, positive and negative changes, percentage changes in populations, and annual rates of change because each captures a different aspect of biodiversity change.

As predicted, we find considerable numerical loss in the avifauna of the EU, although the rate of this decline has slowed and the underlying distribution of species' changes is close to zero, and we find heterogeneity in patterns of change in different bird abundance categories, and in birds associated with different habitats consistent with past studies.

2 | METHODS

2.1 | Data collection and collation

Our analysis covers all breeding bird species native to countries in the EU where adequate data were available (86%; 378 species out of 445 native species that breed in the EU (European Commission, 2018).

2.1.1 | Annual time series

Annual population time series (annual index values and associated standard errors) for 169 common native European bird species...
In line with previous studies (Inger et al., 2015), we expressed all of these. Most population estimates were expressed as numbers. The data cover all 445 native breeding species in the EU. Both surveys to expert opinion; and as a full set of references as possible. Therefore, our results can be thought of as an approximation of the number of breeding individuals of native breeding bird species. The trend and population estimates may be based on (a) complete surveys or statistically robust estimates; (b) extrapolation from limited data; or (c) expert opinion with very limited data (DG Environment, 2017). Approximately 80% of the national population estimates and 70% of the national long-term trends used were based on either option (a) or (b). Data quality varied by country and species, and although including lower quality data may introduce biases of its own, we decided to use all sources of population estimate and population trend data in this study in order for the taxonomic and geographic coverage to be as broad and representative of the full assemblage as possible.

Long-term population trends were used in this study. The period over which the long-term population trend was estimated varied between and within species but was ~1980–2018 in most cases; trend estimates were only included if they covered at least 16 years of this period. Trends were coded as increasing, decreasing, fluctuating around zero, or stable, with quantitative minimum and maximum trend estimates and/or best single value given for the first two categories and for some species/country combinations in the latter two categories. No overall change was assumed to have occurred where national species’ trends were coded as fluctuating, or as stable when no quantitative trend estimate was given.

Annual population time series across the EU were derived from national long-term population trend estimates and population estimates. The proportion of a species’ EU population for which a trend estimate was available varied markedly. We included species for which a trend was available for at least a third of the EU population (N = 356; 80% of all 445 species). We used a lower cutoff for inclusion here compared with the 50% used for EU time series derived from national monitoring schemes (see Section 2.1.1) as the 50% threshold is for the most recent year and the average coverage across the time series is often lower. For each species in each country, we estimated the mean trend ($\bar{T}$; log scale), mean national population estimate ($\bar{E}_n$; log scale), and mean year the population estimate was made ($\overline{y}$). Where a best single value was given, this was used. In the absence of a strict mathematical means to derive standard errors from maximum and minimum values, we roughly approximated standard errors around each mean estimate as a sixth of the difference between the maximum and minimum estimates (covering 99.7% of the distribution if it is assumed to be approximately normal). Where the population estimate maximum and minimum represented a 95% confidence interval, standard errors could be estimated more robustly as a quarter of the difference between the upper and lower estimate of the interval (where a 95% confidence interval is $\pm 1.96$ standard errors above and below the mean). Where maximum and minimum values were absent, the standard error was set to zero.

For each species and each country, we used $\bar{T}$, $\bar{E}_n$, and $\overline{y}$, returning them to the measurement scale, where necessary, to estimate the population size in each year ($y$) 1980–2017 (Equation 1). The resulting country level time series were summed across countries estimates as numbers of individuals, assuming one pair, one male, or one female equaled two individuals.
in each year to obtain an EU-level population time series for each species.

$$\bar{E}_{Ni} = \overline{E}_{1T} \cdot \lambda^{1-(\text{trend period})},$$

where \( \lambda = 1 \text{/(trend period)} \).  \( \text{(1)} \)

We used a bootstrap approach to estimate confidence intervals around the average time series. In each iteration (\( N = 100 \)), the process described earlier was repeated using estimates of \( T, E, \) and \( y \) sampled from a normal distribution described by the average values and standard errors calculated before. The 2.5% and 97.5% quantiles of the bootstrap values for each year were taken as the lower and upper confidence limits.

2.1.3 | Linking annual time series to population estimates

Population size estimates were available for all species in each EU country, from Article 12 reporting. EU-level species’ population estimates were calculated by taking the geometric mean of the summed minimum and maximum national species’ population estimates across countries, where only a best single value was available, and this was treated as both maximum and minimum. As before, we approximated the standard error around these estimates as a sixth of the span of the maximum and minimum values.

The period over which species’ national population estimates were made was approximately 2013–2018 (median start and end years, respectively), but there was variation within and between species. Nevertheless, only 2% of time periods started prior to 2000 and 10% prior to 2010. To obtain a single range estimate per species, we used the median of the country-level start year estimates and the median of the country-level end year estimates. We used these to estimate the midpoint and associated standard error, as described before.

2.1.4 | Covariates

Species were split into four quartiles of abundance, each containing an equal number of species, based upon their EU population estimate and labeled as rare, scarce, common, and abundant. We classified species according to their preferred breeding habitat following BirdLife International (2004), which is based on the habitat association matrix of Tucker and Evans (1997). The nine habitat associations were (1) marine; (2) coastal; (3) inland wetland; (4) tundra, mires and moorland; (5) boreal and temperate forests; (6) Mediterranean forest, shrubland and rocky habitats; (7) agricultural and grassland; and (8) montane grassland. Species that did not fit simply into these categories were labeled as (9) unclassified. A possible alternative description for this last group could be “generalists”; however, we felt that this implied a broad ecological niche, which might not always be the case where species do not fit well into a single one of the habitat classes used. We classified species to a migration strategy as (1) resident, (2) partial migrant, (3) short-distance migrant, and (4) long-distance migrant, following Sanderson et al. (2006). Species were split into four bird groups by family: landbirds, shorebirds, waterbirds, and waterfowl, following Rosenberg et al. (2019).

2.2 | Estimating change in total avian population over time

A single EU-level time series 1980–2017 was selected for each species, using those modeled on multiple national monitoring schemes, where available (\( N = 167 \); 2.1.1); otherwise, those derived from national trend and population estimates (\( N = 211 \); 2.1.2). The time series and the species’ EU population estimates were then analyzed using two different approaches described later, each with different statistical assumptions.

2.2.1 | Bayesian model

First, we used the Bayesian hierarchical model of Rosenberg et al. (2019; Smith, 2019) to estimate change in the abundance of EU birds. The model first creates smoothed species’ time series using a Bayesian GAM for time series modeled on multiple national monitoring schemes and then uses the species time series plus additional data (on breeding habitat and migration strategy) in a hierarchical Bayesian model that models both species- and group-level trends in population size and shrinks uncertain species’ trends toward the group mean. This approach accounts for missing data at the start of species’ indices and incorporates uncertainty in both the annual estimates within the time series, and around the population estimate and the year of the population estimate. The value of initial missing years is set to that of the first year with data, and the variance associated with the missing values is increased by the square of the number of years since nonmissing data. This means that as the number of years between a missing estimate and the closest year with data increases, the estimate has less and less influence on the model output.

2.2.2 | Imputed model

The aforementioned approach shrinks uncertain species’ indices toward the group mean. This could be advantageous, but equally, it could introduce bias if species’ trends correlate with precision, which seems entirely plausible. To verify the earlier results, we used an imputed model like that of Inger et al. (2015), with a bootstrap approach to assess error. As before, the value of initial missing years was set to that of the first year with data. To estimate the average total abundance across species in each year (\( \bar{N} \)), we expressed each species’ time series (\( \bar{N} \)) as a proportion of the value in the average population estimate year (\( \bar{F} \)), multiplied each time series value by the species’ average EU population estimate (\( \bar{F} \)) and summed across species (\( s \)) for each year (\( i \)) (Equation 2).
We used a bootstrap approach \((N = 1000)\) to create confidence intervals. In each iteration, we sampled \(E\) and \(I\) (both log scale), and \(y\) from a normal distribution and estimated the abundance across species in each year as before, after returning values to the measurement scale as necessary. The 2.5% and 97.5% quantiles of the bootstrap values for each year were taken as the lower and upper confidence limits. Our input data for the Imputed model were the same as for the Bayesian model, that is, including the species’ time series generated using the Bayesian GAM (2.2.1).

3 | RESULTS

3.1 | Overall population change in the EU avifauna

We estimate the total number of breeding individuals of native breeding bird species in the EU of the 378 species assessed to have declined by 557m individual birds between 1980 and 2017 \((-17\%; 95\%\) credible interval \(-681\) to \(-433\)) using the Bayesian model and by 623m birds \((-19\%; 95\%\) confidence interval \(-803\) to \(-468\)) using the imputed model (Figure 1a, Table 1). The estimated total abundance in 2017 was 2639 m (2551–2739) using the Bayesian model and 2603 m (2547–2739) using the imputed method.

Visual inspection indicates a difference in the rate of population decline in the late 20th compared with the early 21st century, with much of the decline in bird numbers occurring in the 1980s and 1990s. Piecewise regression on a log scale supports this, indicating a change in slope around the turn of the century (Bayesian: \(\text{slope}_{1980:2001} = -1.00 (-1.04, -0.95)\), \(\text{slope}_{2001:2017} = -0.029 (-0.10, 0.046)\), \(R^2 = 0.99\); imputed: \(\text{slope}_{1980:2000} = -1.01 (-1.07, -0.95)\), \(\text{slope}_{2000:2017} = -0.029 (-0.11, 0.055)\), \(R^2 = 0.99\)), with a decline of approximately 1% per year in the first period and a rate of change not significantly different from zero thereafter.

Given the similarities in the output of the two modeling methods, hereafter, we report only the results of the Bayesian model (Table A1). There were no substantial differences between these findings and the equivalent results based on the imputed model (imputed model outputs are given in Table A2).

Given that some species have increased, the total decline across declining species was 903m and the total increase for increasing species was 341 m (Figure 1b). The distribution of log-transformed species’ population growth rates was positively skewed (skew = 1.14) and leptokurtic (kurtosis = 7.57) (Figure 1c), with the central tendency close to zero (median: 0.00041).

3.2 | Patterns of change by covariate

As predicted, on average rare and scarce bird species in the EU avifauna have less negative population trajectories than more abundant species (Figure 2; Table A1). Overall, rare species showed a 4% decline in abundance as a group and scarce species a 5% decline (Figure 2c: rare \(\%\Delta_{1980-2017} = -4 (-12, 5)\); scarce \(\%\Delta_{1980-2017} = -5 (-10, 2)\)), whereas there was a 25% decline in the total abundance of common species \(\%\Delta_{1980-2017} = -25 (-30, -18)\), and a 17% decline in abundant species \(\%\Delta_{1980-2017} = -17 (-21, -13)\). The median log-transformed population growth rate among rare species was 0.0059 and in scarce species 0.00098 (Figure 2d). In contrast, the average rate of change in common species was −0.0027 and in abundant species −0.0018 (Figure 2d).

The split between abundance categories is correlated with the method of time series derivation, with a higher proportion of rare and scarce species’ time series coming from national trends and...
population estimates than in common or abundant species. However, the trend toward abundant species declining proportionally more than rare species was apparent when all time series derived from national trends and population estimates, where considered separately ($N = 356$, Table A3).

The pattern of change in abundance varies with the species' breeding habitat associations (Figure 3; Table A1). Only five species were associated with montane grassland, so this category was not plotted. We see the largest net decline among species associated with agricultural land and grasslands (Figure 3a; $\Delta_{1980-2017} = -296$ m ($-361$, $-234$)), followed by the "unclassified" species—those species not associated with any single habitat ($\Delta_{1980-2017} = -220$ m ($-325$, $-114$)). When summarized at a species level, the total decline across all declining species was similar between these two groups, but the total increase among species with positive trends was greater in the unclassified group (Figure 3b). The proportional change in the total abundance was greatest for species associated with agricultural land and grasslands ($%\Delta_{1980-2017} = -33$ ($-38$, $-27$)), as well as tundra, mires, and moorland ($%\Delta_{1980-2017} = -28$ ($-36$, $-18$). Species associated with coastal ($\Delta_{1980-2017} = 0.2$ m (0.0, 0.4), $%\Delta_{1980-2017} = 5$ ($-1$, 13)) and Mediterranean habitats ($\Delta_{1980-2017} = 9$ m (1, 20), $%\Delta_{1980-2017} = 23$ (2, 50)) saw small total increases in abundance. In the latter case, this represented a substantial proportional increase, given the low initial abundance of species associated with this habitat. The median log-transformed population growth rate was negative for species associated with agricultural land and grasslands ($-0.52$), boreal and temperate forests ($-0.63$), and tundra, mires, and moorland ($-0.0038$) but positive for those species associated with Mediterranean habitats ($0.012$).

Although resident and long-distance migrant species show similar estimates of total loss over time, the lower total abundance of long-distance migrants means they have declined proportionally more, although the credible intervals overlap (Table A1, long-distance migrants: $\Delta_{1980-2017} = -221$ m ($-270$, $-178$), $%\Delta_{1980-2017} = -33$ ($-39$, $-28$); residents: $\Delta_{1980-2017} = -214$ m ($-312$, $-121$), $%\Delta_{1980-2017} = -21$ ($-29$, $-13$). Among bird groups, Shorebirds show the largest proportional decline (Table A1, $%\Delta_{1980-2017} = -38$ ($-44$, $-32$)), whereas waterfowl show an increase ($%\Delta_{1980-2017} = 23$ (8, 40)).

### Table 1

| Parameter | Bayesian model | Imputed model |
|-----------|----------------|---------------|
|           | Estimate | LCL | UCL | Estimate | LCL | UCL |
| 1980      | 3197    | 3098 | 3300 | 3226    | 3106 | 3451 |
| 2017      | 2639    | 2551 | 2739 | 2603    | 2547 | 2739 |
| Change    | -557    | -681 | -433 | -623    | -803 | -468 |
| % Change  | -17.42  | -21.31 | -13.54 | -19.32  | -24.90 | -14.49 |
| % per annum | -0.52  | -0.63 | -0.40 | -0.58  | -0.77 | -0.42 |

**FIGURE 2** Patterns of change in native breeding bird species in the EU from 1980 to 2017 disaggregated by abundance category. (a) Net change in total abundance (millions of individuals) with 95% credible intervals. (b) Total increase in species with positive trends and total decrease in species with negative trends (millions of individuals). (c) Percent change in total abundance with 95% credible intervals. (d) Box plot of average annual rates of change across species on a log scale. The number of species followed by the range of species' population sizes included in each category is given within brackets.
3.3 | Extreme patterns in species’ change

A small number of species were responsible for a large proportion of the change in numbers observed (Figure 4, species results file available in Burns et al., 2021) in both increasing and decreasing species. *Passer domesticus* (house sparrow) accounts for 27% (247 m) of the total decrease across all declining species. The eight species showing the largest declines account for 69% of the decline across all 175 declining species and the eight species showing the largest increases account for 66% of the increase across all 203 increasing species (Figure 4).

4 | DISCUSSION

Using an extensive dataset and two analytical methods, we demonstrate significant biodiversity loss in the native avifauna of the EU. Specifically, we estimate a decline of 17%–19% in the overall bird abundance since 1980, which equates to a numerical loss of 560–620 m individual birds (Figure 1, Table 1). However, and at first counter-intuitively, the average population growth rate of this assemblage is close to zero, and losses are driven by larger proportional declines in more abundant bird species (Figures 1 and 2). In addition, biodiversity loss is heterogeneous across habitats, with losses pronounced in birds associated with agricultural and grassland habitats, and to a lesser degree, in birds associated with Boreal and temperate forests (Figure 3). Long-distance migrants and shorebirds also appear to have declined more strongly.

Our dataset, although imperfect, represents one of the best datasets of its kind, and we can be reasonably confident in a signal of overall biodiversity loss in the form of a large numerical loss in a bird assemblage at a near continental scale. However, the overall pattern of change conceals variation across species, abundance categories, and habitat. Debate has revolved around the balance between “winners” and “losers” in global biodiversity change (Dornelas et al., 2019; McKinney & Lockwood, 1999), but an equal balance of the two could result in biodiversity loss, or gain, if the magnitude of population changes on either side is not balanced. That is the case in our data-set, where the average population growth rate was close to zero, but the overall population change was negative. This argues, as others have, for a more nuanced view of biodiversity change (Dornelas et al., 2019; Daskalova, 2020; Dornelas et al., 2019; Leung et al., 2020). As in Leung et al. (2020), we also highlight the role of “extreme” population changes in driving the overall pattern of change, both increase and decline (Figure 4). In our case, a broad group of bird species have declined severely for a variety of reasons most closely associated with anthropogenic drivers of land use change (Donald et al., 2001, 2006), so although they are extreme in one sense, they represent a recognized characteristic of this assemblage and are measured with some precision.

Our results are strikingly similar to those of Rosenberg et al. (2019) in North America in showing substantial numerical losses,
heterogeneity in trends among different groups of birds, and “extreme” changes in some species. Rosenberg et al. note that declines in North America parallel studies elsewhere and that the loss of native grassland birds is driven by habitat loss and pesticides, mirroring the loss of farmland birds in Europe (Bowler et al., 2019; Donald et al., 2006). Both studies identify the same groups of birds as of pressing conservation concern, namely, grassland/agricultural birds, shorebirds, and long-distance migrants. Curiously, a small handful of super-abundant species, including *Passer domesticus* (house sparrow) and *Sturnus vulgaris* (starling), drive the numerical decline, although these two species are only native in Europe. The North American dataset starts a decade prior to ours and shows a larger proportional decline in the total abundance. Large losses of farmland birds were reported from 1970 in Europe (Tucker & Heath, 1994); however, these changes varied spatially, with strong declines concentrated among countries in the west of the continent (Donald et al., 2001). Both studies also suggest a slowing in the rate of decline over the last decade. There is evidence that this change might be driven in part by conservation actions that have acted to protect species and create and restore habitats in North America and Europe. In the EU, the Birds Directive (2009/147/EC) and the Habitats Directive (92/43/EEC) provide legal protection to priority species and habitats (European Union, 1992, 2009) and have been shown to benefit target bird species (Donald et al., 2007; Sanderson et al., 2016) and enhance habitat protection (European Environment Agency, 2020). The impact of conservation may explain the trend toward rarer species having more positive annual growth rates (Figure 2). For example, seven of the species in the top decile of growth rates in our dataset are raptors (birds of prey). Many raptor populations have increased in recent decades following increased protection and reductions in pesticides and persecution, as well targeted species’ recovery projects (Deinet et al., 2013; Smart et al., 2010); a pattern mirrored in North America. The Natura 2000 network of protected sites created under the Birds and Habitats Directive has grown rapidly since the early 1990s, rising from 50,000 km² in 1993 to 1,350,000 km² in 2019; nevertheless, only 15% of habitats within the network are in good condition, and the network itself remains incomplete (European Environment Agency, 2020).

Taking our results more broadly, our study supports those reviews that indicate significant biodiversity loss over recent decades (Diaz et al., 2019; Secretariat of the Convention on Biological Diversity, 2020). The degree of loss in EU birds (~17%-19% in nearly 40 years) is similar to the Living Plant Index (LPI) for Europe and Central Asia (~24% average decline in vertebrate trends since 1970, (Almond et al., 2020)), to which many of the species’ time series used here contribute. Other studies describing biological loss in vertebrate populations globally have reported larger declines (Ceballos et al., 2015, 2017). It is argued that historical losses due to land conversion and degradation have left many temperate populations highly depleted, so modern-day trajectories of species in some temperate and tropical comparisons are likely to be different (Newbold et al., 2016). Thus, although we would argue that our dataset is one of the best of its kind available globally, and is a good representation of the EU avifauna, we are not able to generalize these results to other taxa in this region, or to other bird populations and taxa in different parts of the world. Improvements in global monitoring efforts are needed to support similar assessments for different taxonomic groups or regions (Moussy et al., 2021).

The numerical loss of common and abundant species is a concern as it implies damage to the ecosystem structure and function and potentially to the delivery of ecosystem services. Common species may have a lower, higher, or equivalent influence per capita on ecosystem services, but their numerical dominance means that changes in their populations may have large impacts on service provision (Gaston, 2011). For instance, the abundance of common plant species has been seen to strongly influence primary productivity (Smith & Knapp, 2003). Most functional relationships between ecosystem service provision and bird abundance are positive (Gaston et al., 2018). For example, the number of a pest insect species consumed...
increases with increasing bird numbers (Crawford & Jennings, 1989), as does tree seed recruitment in areas of forest regeneration with increased abundance of frugivorous birds (Martínez & García, 2017), and people living in areas of higher bird abundance have lower levels of stress, anxiety, and depression, although effect sizes were low (Cox et al., 2017).

5 | CONCLUSIONS

Here, we demonstrate substantial biodiversity loss using a comprehensive dataset at an assemblage level in the EU. Patterns of change also vary with the abundance class of the species, the habitat they frequent, and other aspects of their ecology, in close parallel to recently described patterns of change in North America. The consistency of patterns of loss across continents makes clear the urgent conservation needs of birds associated with agricultural land/grassland and long-distance migrants. Large declines in the total avian abundance in the EU hide variation in terms of net and gross change in populations, proportional change, and average per annum rates of change. We argue that careful assessment of biodiversity change is needed to understand how to respond in policy terms to the emergent patterns.

Our results support the draft Post-2020 Global Biodiversity Framework of the Convention on Biological Diversity (United Nations Environment Programme, 2021), which calls for increasing conservation efforts to be targeted toward preventing global and national extinctions, and the need to stabilize and restore globally and nationally depleted populations. For the latter to be successful, we need large-scale conservation actions to be implemented widely and effectively across a range of biomes. There is great potential for mechanisms within the EU Biodiversity Strategy 2030 and specifically through the proposed EU ‘restoration law’ to define legally binding targets to restore habitats and species and drive this endeavor. This would require transformative actions across sectors to tackle the nature and climate crises in tandem: protected area networks, species protection, nature-friendly farming, forestry, and fisheries are all key parts of wider society solutions. The datasets used here will play an important role in monitoring the impact of these and related conservation actions.

ACKNOWLEDGEMENTS

We thank the many volunteers who contributed to national bird surveys and the scheme coordinators (Hany Alonso, Marc Anton, Ainars Auniņš, Zoltán Benkő, Mattia Brambilla, Malte Busch, Gianpiero Calvi, Tomasz Chodkiewicz, Przemysław Chylarecki, Jordi Dalmay, Elisabetta de Carli, Juan Carlos del Moral, Antoine Derouaux, Jaanus Derouaux, Tomasz Dąbrowski, Tibor Szép, Guido Tellini Florenzano, Joaquim Teodósio, Norbert Teufelbauer, Sven Trautmann, Tom van der Meij, Chris van Turnhout, Glenn Vermeersch, Zdeněk Vermouzek, Thomas Vikstrøm, Anne Weiserbs) and those who submitted data on bird population sizes and trends across the EU and the national coordinators. PECBMS has been funded by RSPB and the European Commission (Grant Agreement 07.0202/2017/755630/SUB/ENV.D2), and by current service contract No 07.0202/2019/821208/SER/ENV.D.2. The European Red List of Birds project through which data reported by EU Member States under Art. 12 of the Birds Directive was quality checked, validated, and analyzed, and species population sizes compiled, was coordinated by BirdLife International and funded by the European Commission (Service Contract 07.0202/2018/790700/SER/ENV.D.3). We thank Jana Škorpilová, Anne Teller, and Petr Volfíšek for their valuable input to the development and maintenance of PECBMS, Fiona Sanderson for discussion on the study methods, Claire Rutherford for her meticulous work on quality checking the Art. 12 data, and Ariel Brunner for valuable comments on the policy relevance.

CONFLICT OF INTEREST

None declared.

AUTHOR CONTRIBUTIONS

Fiona Burns: Conceptualization (equal); Formal analysis (lead); Methodology (lead); Writing-original draft (equal); Writing-review & editing (equal). Mark A. Eaton: Conceptualization (equal); Methodology (supporting); Writing-review & editing (equal). Ian J. Burfield: Data curation (equal); Methodology (supporting); Writing-review & editing (equal). Alena Kvaňová: Data curation (equal); Methodology (supporting); Writing-review & editing (equal). Eva Šilarová: Data curation (equal); Methodology (supporting); Writing-review & editing (equal). Anna Staneva: Data curation (equal); Methodology (supporting); Writing-review & editing (equal). Richard D. Gregory: Conceptualization (equal); Methodology (supporting); Writing-original draft (equal); Writing-review & editing (equal).

DATA AVAILABILITY STATEMENT

The two input datasets and the species’ level results have been archived on Zenodo (Burns et al., 2021).

ORCID

Fiona Burns https://orcid.org/0000-0002-5362-5552

REFERENCES

Almon, R., Grooten, M., & Peterson, T. (2020). Living planet report 2020-bending the curve of biodiversity loss. World Wildlife Fund. Bowler, D. E., Heldbjerg, H., Fox, A. D., de Jong, M., & Böehning-Gaese, K. (2019). Long-term declines of European insectivorous bird
populations and potential causes. Conservation Biology, 33, 1120-1130. https://doi.org/10.1111/cobi.13307

Brlík, V., Šilarová, A., Škorpilová, J., Alonso, H., Anton, M., Aunins, A., Benkó, Z., Biver, G., Busch, M., Chodkiewicz, T., Chylarecki, P., Coomes, D., de Carli, E., del Morai, J. C., Derouaux, A., Escandell, V., Eskildsen, D. P., Fontaine, B., Poppen, R. P. B., ... Klvaňová, A. (2020). Data from: Long-term and large-scale multispecies dataset tracking population changes of common European breeding birds. https://doi.org/10.5281/zenodo.4590199

Brlík, V., Šilarová, A., Škorpilová, J., Alonso, H., Anton, M., Aunins, A., Benkó, Z., Biver, G., Busch, M., Chodkiewicz, T., Chylarecki, P., Coomes, D., de Carli, E., del Morai, J. C., Derouaux, A., Escandell, V., Eskildsen, D. P., Fontaine, B., Poppen, R. P. B., ... Klvaňová, A. (2021). Long-term and large-scale multispecies dataset tracking population changes of common European breeding birds. Scientific Data, 8, 1–9. https://doi.org/10.1038/s41597-021-00804-2

Buckland, S., & Johnston, A. (2017). Monitoring the biodiversity of regions: Key principles and possible pitfalls. Biological Conservation, 214, 23–34. https://doi.org/10.1016/j.biocon.2017.07.034

Burns, F., Eaton, M. A., Burfield, I. J., Klvaňová, A., Šilarová, E., Staneva, A., & Gregory, R. D. (2021). Data from: Abundance decline in the avifauna of the European Union reveals global similarities in biodiversity change: Input datasets & species results. Zenodo. https://doi.org/10.5281/zenodo.5544548

Ceballos, G., Ehrlich, P. R., Barnosky, A. D., García, A., Pringle, R. M., & Palmer, T. M. (2015). Accelerated modern human–induced species losses: Entering the sixth mass extinction. Science Advances, 1, e1400253. https://doi.org/10.1126/sciadv.1400253

Ceballos, G., Ehrlich, P. R., & Dirzo, R. (2017). Biological annihilation via the ongoing sixth mass extinction signaled by vertebrate population losses and declines. Proceedings of the National Academy of Sciences of the United States of America, 114, E6089–E6096. https://doi.org/10.1073/pnas.170494114

Cox, D. T. C., Shanahan, D. F., Hudson, H. L., Plummer, K. E., Siriwardena, G. M., Fuller, R. A., Anderson, K., Hancock, S., & Gaston, K. J. (2017). Doses of neighborhood nature: the benefits for mental health of living with nature. BioScience, 67, 147–155. https://doi.org/10.1093/biosci/biw173

Crawford, H. S., & Jennings, D. T. (1989). Predation by birds on spruce budworm Choristoneura fumiferana: functional, numerical, and total responses. Ecology, 70, 152–163. https://doi.org/10.2307/1938422

Deinet, S., Jeronymidou, C., McMae, L., Burfield, I. J., Foppen, R. P. C., Collen, B., & Böhm, M. (2013). Wildlife comeback in Europe: The recovery of selected mammal and bird species. Final report to Rewilding Europe by ZSL. BirdLife International and the European Bird Census Council. ZSL.

DG Environment (2017). Reporting under Article 12 of the Birds Directive: Explanatory notes and guidelines for the period 2013–2018. Brussels. http://cdr.eionet.europa.eu/help/birds_art12

Díaz, S., Settele, J., Brondizio, E. S., Ngo, H. T., Agard, J., Arnessh, A., Balvanera, P., Brauman, K. A., Butchart, S. H. M., Chan, K. M. A., Garibaldi, L. A., Ichii, K., Liu, J., Subramanian, S. M., Midgley, G. F., Miloslavich, P., Molnár, Z., Obura, D., Pfaff, A., ... Zayas, C. N. (2019). Pervasive human-driven decline of life on Earth points to the need for transformative change. Science, 366. https://doi.org/10.1126/science.aax3100

Donald, P. F., Green, R., & Heath, M. (2001). Agricultural intensification and the collapse of Europe’s farmland bird populations. Proceedings of the Royal Society of London. Series B: Biological Sciences, 268, 25–29.

Donald, P. F., Sanderson, F. J., Burfield, I. J., Bierman, S. M., Gregory, R. D., & Walczyk, Z. (2007). International conservation policy delivers benefits for birds in Europe. Science, 317, 810–813. https://doi.org/10.1126/science.1146002

Donald, P. F., Sanderson, F. J., Burfield, I. J., & Van Bommel, F. P. (2006). Further evidence of continent-wide impacts of agricultural intensification on European farmland birds, 1990–2000. Agriculture, Ecosystems & Environment, 116, 189–196. https://doi.org/10.1016/j.agee.2006.02.007

Dornelas, M., & Daskalova, G. N. (2020). Nuanced changes in insect abundance. Science, 368, 368–369. https://doi.org/10.1126/science.abb6861

Dornelas, M., Gotelli, N. J., Shimadzu, H., Moyes, F., Magurran, A. E., & McGill, B. J. (2019). A balance of winners and losers in the Anthropocene. Ecology Letters, 22, 847–854. https://doi.org/10.1111/ele.13242.

EBCC/RSBP/BirdLife/CSO (2020). EU species indices and trends till 2017. PECBMS.

Eionet (2020). Article 12 web tool: population status and trends of birds under Article 12 of the Birds Directive. European Environment Agency. https://nature-art12.eionet.europa.eu/article12/

European Commission (2018). List of the birds of the European Union. European Commission, N2K Group and BirdLife International.

European Environment Agency (2020). State of nature in the EU: Results from reporting under the nature directives 2013-2018. EEA Report No 10/2020. ISSN 1725-9177. https://www.eea.europa.eu/publications/state-of-nature-in-the-eu-2020

European Union (1992). Directive 92/43/EEC on the conservation of natural habitats and of wild fauna and flora of May1992. http://www.eurex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:31992L0043&from=EN

European Union (2009). Directive 2009/147/EC of the European parliament and of the council. http://eurlex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32009L0147&from=EN

Gaston, K. J. (2010). Valuing common species. Science, 327, 154–155. https://doi.org/10.1126/science.1182818

Gaston, K. J. (2011). Common ecology. BioScience, 61, 354–362. https://doi.org/10.1525/bios.2011.61.5.4.

Gaston, K. J., Cox, D. T. C., Canavelli, S. B., García, D., Hughes, B., Maas, B., Martinez, D., Ogada, D., & Inger, R. (2018). Population abundance and ecosystem service provision: The case of birds. BioScience, 68, 264–272. https://doi.org/10.1093/biosci/biy005

Gaston, K. J., & Fuller, R. A. (2008). Commonness, population depletion and conservation biology. Trends in Ecology & Evolution, 23, 14–19. https://doi.org/10.2326/9781839000079

Gregory, R. D., Skorpilova, J., Vorisek, P., & Butler, S. (2019). An analysis of trends, uncertainty and species selection shows contrasting trends of widespread forest and farmland birds in Europe. Ecological Indicators, 103, 676–687. https://doi.org/10.1016/j.ecolind.2019.04.064

Gonzalez, A., Cardinale, B. J., Allington, G. R. H., Byrnes, J., Arthur Endsley, K., Brown, D. G., Hooper, D. U., Isbell, F., O’Connor, M. I., & Loreau, M. (2016). Estimating local biodiversity change: a critique of papers claiming no net loss of local diversity. Ecology, 97, 1949–1960. https://doi.org/10.1890/15-1759.1

Gregory, R. D., & van Strien, A. (2010). Wild bird indicators: using composite population trends of birds as measures of environmental health. Ornithological Science, 9, 3–22. https://doi.org/10.2326/09s.9.3

Gregory, R. D., van Strien, A., Vorisek, P., Gmelig Meyling, A. W., Noble, D. G., Foppen, R. P. B., & Gibbons, D. W. (2005). Developing indicators for European birds. Philosophical Transactions of the Royal Society B: Biological Sciences, 360, 269–288. https://doi.org/10.1098/rstb.2004.1602

Harris, S. J., Massimino, D., Balmer, D. E., Eaton, M. A., Noble, D. G., Pearce-Higgins, J. W., Woodcock, P., & Gillings, S. (2020). The Breeding Bird Survey 2019. BTO Research Report 726. Thetford. www.bto.org/bbs
Inger, R., Gregory, R., Duffy, J. P., Stott, I., Voirišek, P., & Gaston, K. J. (2015). Common European birds are declining rapidly while less abundant species’ numbers are rising. Ecology Letters, 18, 28–36. https://doi.org/10.1111/ele.12387

Jörg-Hickfang, T., Hofmann, M., Martins, I., Mimet, A., McGill, B., & Pereira, H. M. (2020). Letter to decline: Response to: Decline of the North American avifauna. Science. https://doi.org/10.1126/science.aaw1313

Keller, B., Martínez, D., & García, D. (2017). Role of avian seed dispersers in tree re-convulsion. Global Change Biology, 3948–3959. https://doi.org/10.1111/gcb.13292

Rosenberg, K. V., Dokter, A. M., Blancher, P. J., Sauer, J. R., Smith, A. C., & Knapp, A. K. (2003). Dominant species maintain ecosystem function with non-random species loss. Ecology Letters, 6, 509–517. https://doi.org/10.1046/j.1461-0248.2003.00454.x

Tucker, G. M., Evans, M. J. (1997). Habitats for birds in Europe: A conservation strategy for the wider environment (Vol. 6). BirdLife International.

Tucker, G., & Heath, M. (1994). Birds in Europe: Their conservation status. Birdlife International (BirdLife Conservation Series 3).

United Nations Environment Programme. (2021). Convention on biological diversity. First draft of the post-2020 global biodiversity framework, CBD/WG2020/3/3. https://www.cbd.int/doc/c/abb5/591f/2e46096d3f0330b08ce87a45/wg2020-03-03-en.pdf

Vellend, M., Baeten, L., Myers-Smith, I. H., Elmundorff, S. C., Beauséjour, R., Brown, C. D., De Frenne, P., Verheyen, K., & Wipf, S. (2013). Global meta-analysis reveals no net change in local-scale plant biodiversity over time. Proceedings of the National Academy of Sciences of the United States of America, 110, 19456–19459. https://doi.org/10.1073/pnas.1312779110

Vellend, M., Dornelas, M., Baeten, L., Beauséjour, R., Brown, C. D., De Frenne, P., Elmundorff, S. C., Gotelli, N. J., Moyes, F., Myers-Smith, I. H., Magurran, A. E., McGill, B. J., Shimadzu, H., & Sievers, C. (2017). Estimates of local biodiversity change over time stand up to scrutiny. Ecology, 98, 583–590. https://doi.org/10.1002/ecy.1660

How to cite this article: Burns, F., Eaton, M. A., Burfield, I. J., Kvaňová, A., Šťarlová, E., Staneva, A., & Gregory, R. D. (2021). Abundance decline in the avifauna of the European Union reveals cross-continental similarities in biodiversity change. Ecology and Evolution, 11, 16647-16660. https://doi.org/10.1002/ece3.8282
TABLE A1  Total change (millions of individuals) and proportional change in breeding bird numbers in the EU, estimated using the Bayesian hierarchical model (see Section 2.2.1) and showing the median estimate and the lower and upper bounds of the 95% credible interval in each case, disaggregated by abundance category, bird group, migration strategy and breeding habitat

| Grouping            | Category          | N  | Total change (m) | Proportional change |
|---------------------|-------------------|----|------------------|---------------------|
|                     |                   |    | Median   | LCI      | UCI      | Median   | LCI      | UCI      |
| All                 |                   | 378| −557.01  | −681.47  | −432.76  | −0.174   | −0.213   | −0.135   |
| Abundance category  | Abundant          | 95 | −524.11  | −647.53  | −398.89  | −0.171   | −0.207   | −0.133   |
|                     | Common            | 95 | −33.02   | −40.30   | −23.00   | −0.252   | −0.300   | −0.176   |
|                     | Scarce            | 94 | −0.44    | −1.02    | 0.21     | −0.046   | −0.102   | 0.022    |
|                     | Rare              | 94 | −0.03    | −0.12    | 0.04     | −0.039   | −0.119   | 0.048    |
| Bird group          | Landbirds         | 255| −548.65  | −673.22  | −424.34  | −0.174   | −0.209   | −0.137   |
|                     | Shorebirds        | 31 | −6.13    | −7.29    | −4.95    | −0.384   | −0.441   | −0.321   |
|                     | Waterbirds        | 64 | −3.69    | −5.35    | −1.95    | −0.165   | −0.227   | −0.089   |
|                     | Waterfowl         | 28 | 1.60     | 0.59     | 2.83     | 0.226    | 0.080    | 0.402    |
| Migration strategy  | Long-distance     | 115| −221.17  | −269.57  | −178.29  | −0.332   | −0.385   | −0.279   |
|                     | Partial migrant   | 127| −116.57  | −182.24  | −49.83   | −0.078   | −0.120   | −0.034   |
|                     | within Europe     | 81 | −213.65  | −312.15  | −120.93  | −0.213   | −0.293   | −0.125   |
|                     | Resident          | 55 | −4.15    | −8.30    | 0.28     | −0.118   | −0.225   | 0.008    |
|                     | Short-distance    |    |          |          |          |          |          |          |
|                     | migrant           |    |          |          |          |          |          |          |
| Breeding habitat    | Agricultural and | 54 | −295.90  | −360.57  | −233.62  | −0.327   | −0.381   | −0.268   |
|                     | grassland         | 70 | −42.15   | −63.60   | −20.74   | −0.089   | −0.132   | −0.044   |
|                     | Boreal and        | 21 | 0.17     | −0.04    | 0.38     | 0.055    | −0.014   | 0.129    |
|                     | temperate forests |    |          |          |          |          |          |          |
|                     | Coastal           | 82 | −219.61  | −324.91  | −114.25  | −0.129   | −0.185   | −0.069   |
|                     | Unclassified      | 51 | −7.05    | −10.82   | −2.86    | −0.134   | −0.201   | −0.055   |
|                     | Inland wetlands   | 26 | 0.19     | −0.84    | 1.46     | 0.024    | −0.092   | 0.174    |
|                     | Marine            | 33 | 9.37     | 0.88     | 19.66    | 0.233    | 0.020    | 0.503    |
|                     | Mediterranean     |    |          |          |          |          |          |          |
|                     | habitats          |    |          |          |          |          |          |          |
|                     | Montane grassland | 5  | 0.46     | 0.27     | 0.71     | 0.646    | 0.360    | 1.001    |
|                     | Tundra, mires,    | 34 | −1.90    | −2.61    | −1.18    | −0.275   | −0.362   | −0.175   |
|                     | and moorland      |    |          |          |          |          |          |          |
|                     | Unknown           | 2  | 0.001    | −0.0005  | 0.003    | 0.318    | −0.128   | 0.961    |
| Grouping   | Category          | N  | Total change (m) | Proportional change |
|-----------|-------------------|----|------------------|---------------------|
|           |                   |    | Estimate         | LCI                 | UCI                 | Estimate | LCI   | UCI   |
| All       |                   | 378| -623.18          | -803.24            | -467.52            | -19.32   | -24.90| -14.49|
| Abundance |                   |    |                  |                     |                     |          |       |       |
| category  |                   |    |                  |                     |                     |          |       |       |
|           | Abundant          | 95 | -583.95          | -769.75            | -431.16            | -18.91   | -24.92| -13.96|
|           | Common            | 95 | -38.79           | -70.98             | -0.96              | -30.23   | -55.32| -0.75 |
|           | Scarce            | 94 | 0.45             | -0.96              | 0.14               | -5.49    | -11.66| 1.74  |
|           | Rare              | 94 | 0.01             | -0.06              | 0.05               | 1.00     | -8.62 | 6.92  |
| Bird group|                   |    |                  |                     |                     |          |       |       |
|           |                   |    |                  |                     |                     |          |       |       |
|           | Landbirds         | 255| -613.58          | -804.77            | -450.84            | -19.30   | -25.31| -14.18|
|           | Shorebirds        | 31 | -6.39            | -7.83              | -5.20              | -39.81   | -48.78| -32.39|
|           | Waterbirds        | 64 | -4.87            | -7.22              | -2.92              | -20.88   | -30.95| -12.49|
|           | Waterfowl         | 28 | 1.66             | 0.75               | 2.67               | 23.51    | 10.60 | 37.79 |
| Migration |                   |    |                  |                     |                     |          |       |       |
| strategy  |                   |    |                  |                     |                     |          |       |       |
|           | Long-distance     | 115| -255.75          | -374.05            | -182.55            | -36.99   | -54.11| -26.41|
|           | Partial migrant   | 127| -133.46          | -205.99            | -65.63             | -8.97    | -13.85| -4.41 |
|           | within Europe     | 127| -133.46          | -205.99            | -65.63             | -8.97    | -13.85| -4.41 |
|           | Resident          | 81 | -228.74          | -357.64            | -124.08            | -22.61   | -35.36| -12.27|
|           | Short-distance    | 55 | -5.23            | -10.85             | -0.36              | -14.53   | -30.14| -1.01 |
| Breeding  |                   |    |                  |                     |                     |          |       |       |
| habitat   |                   |    |                  |                     |                     |          |       |       |
|           | Agricultural and  | 54 | -331.36          | -443.21            | -249.87            | -35.65   | -47.69| -26.89|
|           | grassland         |    |                  |                     |                     |          |       |       |
|           | Boreal and        | 70 | -48.07           | -70.19             | -26.27             | -10.09   | -14.74| -5.52 |
|           | temperate forests |    |                  |                     |                     |          |       |       |
|           | Coastal           | 21 | 0.13             | -0.07              | 0.30               | 4.34     | -2.45 | 9.77  |
|           | Unclassified      | 82 | -242.22          | -376.09            | -111.30            | -14.18   | -22.02| -6.52 |
|           | Inland wetlands   | 51 | -8.12            | -12.66             | -3.32              | -15.39   | -24.00| -6.30 |
|           | Marine            | 26 | -0.31            | -1.36              | 0.47               | -3.55    | -15.87| 5.43  |
|           | Mediterranean     | 33 | 8.26             | -0.86              | 19.80              | 20.47    | -2.14 | 49.09 |
|           | habitats          |    |                  |                     |                     |          |       |       |
|           | Montane grassland | 5  | 0.47             | 0.35               | 0.58               | 65.88    | 48.88 | 81.27 |
|           | Tundra, mires,    | 34 | -1.97            | -2.67              | -1.29              | -28.56   | -38.73| -18.76|
|           | and moorland      |    |                  |                     |                     |          |       |       |
|           | Unknown           | 2  | 0.001            | 0.0002             | 0.002              | 29.83    | 4.71  | 56.46 |
| Year | Rare  
(N = 88; 5–16 k) | | | Scarce  
(N = 89; 16 k–127 k) | | | Common  
(N = 89; 128 k–1.5 m) | | | Abundant  
(N = 90; 1.5–222 m) | |
|------|---------------|---|---|----------------|---|---|---------------|---|---|---------------|---|---|
|      |  Est | LCL  | UCL  |  Est | LCL  | UCL  |  Est | LCL  | UCL  |  Est | LCL  | UCL  |  |
| 1980 | 0.56 | 0.52  | 0.75  | 5.41 | 5.13  | 6.56  | 59.67 | 57.99 | 62.81 | 1,765.0 | 1,748.3 | 1,802.2 |
| 2017 | 0.53 | 0.52  | 0.60  | 5.16 | 5.07  | 5.80  | 48.70 | 48.10 | 50.10 | 1,550.4 | 1,535.8 | 1,579.3 |
| Change (m) | −0.02 | −0.16 | 0.004 | −0.25 | −1.17 | 0.03 | −10.97 | −14.03 | −8.93 | −214.68 | −230.75 | −202.11 |
| Change (%) | −4.29 | −28.02 | 0.74 | −4.55 | −21.69 | 0.62 | −18.38 | −23.52 | −14.96 | −12.16 | −13.07 | −11.45 |