

Extended Data Fig. 1 | Conceptual diagrams showing directional patterns of long-distance seed dispersal by migratory birds and phenological overlaps between seed-dispersal periods and bird migrations. a, Yellow and black arrows denote long-distance seed dispersal within and beyond the current range of a plant species, respectively. Seed dispersal mediated by birds migrating south (left), non-migrating birds (centre) and birds migrating north (right). The colour gradient from red to blue represents a climatic gradient from warmer to cooler latitudes (from south to north in the Northern Hemisphere), respectively. In the diagram on the right, seed dispersal within the range is necessary for warm-adapted populations to colonize cooler areas that are warming owing to climate change, whereas seed dispersal beyond the range is necessary for range shifts. **b**, Three hypothetical examples of phenological overlap between the seed-dispersal period of plant species *i* and bird species *j* while the bird migrates northwards (top), southwards (middle) or during both migrations (bottom). The examples include a wintering migrant with a winter-spring fruiting plant (top); a summer migrant with a summer-autumn fruiting plant (middle); and a transient migrant with an autumn-winter fruiting plant (bottom). In some cases, there is also phenological overlap during non-migration periods. More details on phenological overlaps in relation to the migratory strategy of birds are provided in Supplementary Fig. 3.



Extended Data Fig. 2 | Estimated interaction frequencies of plant species within each study network with birds migrating northwards, southwards or not migrating. Blue, interactions during northward migration; red, interactions during southward migration; grey, non-migration interactions.

Each panel represents a seed-dispersal network. The left column of panels includes Mediterranean networks, whereas the right column includes temperate networks. DE, Germany; ES, Spain; IT, Italy; PL, Poland; PT, Portugal; UK, United Kingdom.



Million years before present

Extended Data Fig. 3 | **Variables of the seed-dispersal phenology across the phylogenetic tree of plants.** Phylogenetic signal was tested in plant-species means across networks in start and end dates (D_{start} and D_{end}), as well as in length ($D_{length} = D_{end} - D_{start}$) of the seed-dispersal period (n = 81 plant species) by means of Pagels' λ , as described in 'Phylogenetic signal in plants' in 'Statistical analyses' (Methods). The three phenological variables showed significant phylogenetic signal (D_{start} , $\lambda = 0.800$, P = 0.0103; D_{end} , $\lambda = 0.781$, P = 0.0015; and D_{length} , $\lambda = 0.419$, P = 0.0343). To test for phylogenetic signal, we previously calculated the species-level means for D_{start} , D_{end} and D_{length} across bioclimates (Extended Data Fig. 6). For this reason, we assessed the amount of variance in these phenological variables that is accounted for by bioclimate, as compared to that accounted for by species through linear-mixed models (LMMs) that included 'bioclimate' as fixed factor and 'plant species' as random factor to account for the repeated measures per species. Bioclimate accounted for only a minor fraction of variance (1–3%) in D_{startv} , D_{end} and D_{length} , as shown by the marginal R^2 values (variance explained by fixed effects; $R^2_{\text{LMM}(m)} = 0.028, 0.01$ and 0.023, respectively). By contrast, the high conditional R^2 values (variance explained by both fixed and random effects; $R^2_{\text{LMM}(m)} = 0.780, 0.845$ and 0.643, respectively) indicated that plant species accounted for most variance in the three phenological variables. LMMs were fitted with the R package package lme4 (v.1.1-19)¹⁰⁵.



Extended Data Fig. 4 | Number of migratory bird species interacting with plants during migration per network in relation to migration direction and **biome.** Large dots and bars denote means \pm 95% confidence intervals estimated by a GLMM, whereas circles denote values for each seed-dispersal network (n = 26 observations, 13 networks × 2 directions). Only migration direction had significant effects on the number of migratory bird species interacting with plants during migration in the GLMM (Poisson family and log-link function) testing the effects of migration direction (Wald $\chi^2 = 11.08$, P=0.0009), biome (Wald $\chi^2=0.17$, P=0.6789) and their interaction (Wald $\chi^2 = 0.02, P = 0.8921$). Model estimates ± s.e.: intercept = 2.297 ± 0.156; direction $(northward) = -0.500 \pm 0.208; biome (temperate) = -0.091 \pm 0.215; direction$ $(northward) \times biome(temperate) = 0.039 \pm 0.288; southward and$ Mediterranean were used as the reference categories (intercepts) for the factors direction and biome, respectively. A mean of 9.5 bird species per community dispersed plants during their southward migration, but only 5.9 species did so during the northward migration.





temperate biome (τ =0.588, P=0.0006), indicating that bird species generally display a proportional role in both migrations. However, the cumulative species strength in the Mediterranean and temperate biome were not correlated, neither across the northward (τ =0.276, P=0.1089) nor across the southward subnetworks (τ =0.263, P=0.0764) (correlation between left and right panels in **a**). These results indicate discordance between biomes in the identity of bird species contributions to community-wide seed dispersal during each migration. Pearson's *r* yielded qualitatively similar results, with higher coefficient values in the significant correlations (r=0.946 and 0.847).



Extended Data Fig. 6 | Bioclimate-level plant phenology from several sources. Subset of 16 out of the 81 plant species present in the study networks illustrating how, in many cases, we obtained data on seed-dispersal phenology

from several sources for the same plant species-bioclimate combination. Colour codes denote different data sources. A vertical grey line divides the calendar year.

Extended Data Table 1 | Characteristics of the European seed-dispersal networks that we studied

Ν	Country	Network name	Source	Sampling type	Biome	Bioclimate	Latitude	Longitude	Years (<i>n</i>)	N _{plant}	N _{bird}	N _{int}
1	Spain	Hato Ratón	А	Mist-netting	Mediterranean	thermo-	37.1804	-6.3240	1981–1983 (2)	16	17	120
2	Spain	Nava Correhuelas	А	Observations	Mediterranean	supra-	37.9409	-2.7927	1997–1999 (2)	22	21	111
3	Spain	Garrapilos	В	DNA-barcoding	Mediterranean	thermo-	36.6589	-5.9493	2013–2015 (2)	14	21	56
4	Spain	Cabañeros	в	DNA-barcoding	Mediterranean	meso-	39.3213	-4.2896	2016–2017 (1)	16	14	44
5	Spain	Arbazal	в	DNA-barcoding	temperate	thermo/meso-	43.4313	-5.4971	2016–2017 (1)	14	14	52
6	Portugal	Vale Soeiro	С	Mist-netting	Mediterranean	meso-	40.3127	-8.4035	2012–2018 (6)	21	13	76
7	Italy	Ficuzza	В	DNA-barcoding	Mediterranean	meso-	37.8923	13.3749	2016–2017 (1)	13	12	30
8	UK	Buckinghamshire	D	Observations	temperate	meso/supra-	51.8910	-0.9120	1980–1985 (5)	29	19	204
9	UK	Wytham Woods	Е	Observations	temperate	meso/supra-	51.7667	-1.3333	1979–1980 (1)	8	8	24
10	UK	Bradfield Woods	В	DNA-barcoding	temperate	meso/supra-	52.1808	0.8239	2016–2017 (1)	15	11	36
11	Germany	Hesse Highlands	F	Observations	temperate	supra-	51.3957	8.9427	1997–1999 (2)	28	18	128
12	Germany	Bauerbach	В	DNA-barcoding	temperate	supra-	50.7950	8.8230	2016–2017 (1)	10	9	30
13	Poland	Hebdów	в	DNA-barcoding	temperate	supra-	50.1429	20.4274	2016–2017 (1)	11	16	38

*N*_{plant}, *N*_{bird} and *N*_{int} denote the number of plant species, bird species and plant-bird interactions in each network, respectively. Network biomes were obtained from refs. ^{47,48}; network bioclimates were obtained from ref. ⁴⁸. Source A is ref. ⁴⁹; source B is this study (EU project MobileLinks); source C is this study (unpublished data provided by L.P.d.S. and R.H.H.); source D is ref. ⁵⁰; source E is ref. ⁵¹.

Extended Data Table 2 | List of bird and plant species of the 13 study networks

Bird species	list	Plant species list				
Bird species	Bird family	Plant species	Plant family	Plant species	Plant family	
Alectoris rufa	Phasianidae	Amelanchier lamarckii	Rosaceae	Prunus avium	Rosaceae	
Columba palumbus	Columbidae	Amelanchier ovalis	Rosaceae	Prunus domestica	Rosaceae	
Corvus corax	Corvidae	Arbutus unedo	Ericaceae	Prunus mahaleb	Rosaceae	
Corvus corone	Corvidae	Arum italicum†	Araceae	Prunus padus	Rosaceae	
Corvus monedula	Corvidae	Arum maculatum†	Araceae	Prunus prostrata	Rosaceae	
Cyanistes caeruleus	Paridae	Asparagus acutifolius	Asparagaceae	Prunus serotina	Rosaceae	
Cyanopica cooki	Corvidae	Asparagus aphyllus	Asparagaceae	Prunus spinosa	Rosaceae	
Dendrocopos major	Picidae	Berberis vulgaris	Berberidaceae	Pyrus amygdaliformis	Rosaceae	
Emberiza calandra	Emberizidae	Bryonia dioica†	Cucurbitaceae	Pyrus bourgaeana*	Rosaceae	
Erithacus rubecula	Muscicapidae	Cornus sanguinea	Cornaceae	Rhamnus alaternus	Rhamnaceae	
Falco tinnunculus	Falconidae	Cotoneaster granatensis	Rosaceae	Rhamnus cathartica	Rhamnaceae	
Ficedula hypoleuca	Muscicapidae	Crataegus monogyna	Rosaceae	Rhamnus lycioides	Rhamnaceae	
Fringilla coelebs	Fringillidae	Daphne gnidium	Thymelaeaceae	Rhamnus saxatilis	Rhamnaceae	
Gallinula chloropus	Rallidae	Daphne laureola	Thymelaeaceae	Ribes rubrum	Grossulariaceae	
Garrulus glandarius	Corvidae	Euonymus europaeus	Celastraceae	Rosa canina	Rosaceae	
Lanius excubitor	Laniidae	Ficus carica	Moraceae	Rosa sempervirens	Rosaceae	
Luscinia megarhynchos	Muscicapidae	Fragaria vesca†	Rosaceae	Rubia peregrina‡	Rubiaceae	
Muscicapa striata	Muscicapidae	Frangula alnus	Rhamnaceae	Rubus fruticosus	Rosaceae	
Oriolus oriolus	Oriolidae	Hedera helix‡	Araliaceae	Rubus idaeus	Rosaceae	
Parus major	Paridae	Hedera hibernica‡	Araliaceae	Rubus ulmifolius	Rosaceae	
Phasianus colchicus	Phasianidae	llex aquifolium	Aquifoliaceae	Ruscus aculeatus	Asparagaceae	
Phoenicurus ochruros	Muscicapidae	Jasminum fruticans	Oleaceae	Sambucus nigra	Adoxaceae	
Phoenicurus phoenicurus	Muscicapidae	Juniperus communis	Cupressaceae	Sambucus racemosa	Adoxaceae	
Pica pica	Corvidae	Juniperus oxycedrus	Cupressaceae	Smilax aspera‡	Smilacaceae	
Picus sharpei	Picidae	Juniperus phoenicea	Cupressaceae	Solanum dulcamara†	Solanaceae	
Picus viridis	Picidae	Juniperus sabina	Cupressaceae	Solanum nigrum†	Solanaceae	
Saxicola torquatus	Muscicapidae	Ligustrum vulgare	Oleaceae	Sorbus aria	Rosaceae	
Sitta europaea	Sittidae	Lonicera arborea	Caprifoliaceae	Sorbus aucuparia	Rosaceae	
Streptopelia decaocto	Columbidae	Lonicera caprifolium‡	Caprifoliaceae	Sorbus torminalis	Rosaceae	
Sturnus unicolor	Sturnidae	Lonicera etrusca‡	Caprifoliaceae	Symphoricarpos albus	Caprifoliaceae	
Sturnus vulgaris	Sturnidae	Lonicera periclymenum‡	Caprifoliaceae	Dioscorea communis†	Dioscoreaceae	
Sylvia atricapilla	Sylviidae	Lonicera xylosteum	Caprifoliaceae	Taxus baccata	Taxaceae	
Sylvia borin	Sylviidae	Malus sylvestris	Rosaceae	Viburnum lantana	Adoxaceae	
Sylvia cantillans	Sylviidae	Morus alba	Moraceae	Viburnum opulus	Adoxaceae	
Sylvia communis	Sylviidae	Morus nigra	Moraceae	Viburnum tinus	Adoxaceae	
Sylvia conspicillata	Sylviidae	Myrtus communis	Myrtaceae	Viscum album	Santalaceae	
Sylvia curruca	Sylviidae	Olea europaea	Oleaceae	Vitis vinifera‡	Vitaceae	
Sylvia hortensis	Sylviidae	Osyris alba	Santalaceae			
Sylvia melanocephala	Sylviidae	Phillyrea angustifolia	Oleaceae			
Sylvia undata	Sylviidae	Phillyrea latifolia	Oleaceae			
Turdus iliacus	Turdidae	Phytolacca americana†	Phytolaccaceae			
Turdus merula	Turdidae	Pistacia lentiscus	Anacardiaceae			
Turdus philomelos	Turdidae	Pistacia terebinthus	Anacardiaceae			
Turdus pilaris	Turdidae	Polygonatum odoratum†	Asparagaceae			
Turdus torquatus	Turdidae					
Turdus viscivorus	Turdidae					

We followed taxonomy from 'Birds of the World' (www.birdsoftheworld.org)⁷⁸ for birds and a previously published⁹⁶ phylogenetic tree (ALLMB) for plants. Plants are defined as herbs (†), woody vines (†) or trees and shrubs (all other species).

*Pyrus bourgaeana (Iberian wild pear) was not present in ref. ⁹⁶ but 'World Flora Online' (www.worldfloraonline.org) considers this species as a synonym of Pyrus communis auct. iber. We thus matched P. bourgaeana to P. communis in the phylogenetic tree to test for phylogenetic signal.

Extended Data Table 3 | Significance of the fixed factors migration direction and biome, and their interaction, in GLMMs testing effects on seed-dispersal interactions of plants with migrating birds

Fixed-effects	(<i>i</i>) Propo sr (Binon	rtion of plant pecies nial, logit link)	(<i>ii</i>) Frequency of seed-dispersal interactions (Beta, logit link)		(<i>iii</i>) Number of bird species per plant (Poisson, log link)	
Hypothesis testing	χ²	Р	X ²	Р	χ²	Р
Direction (D)	51.02	2.0 × 10 ⁻¹⁶	159.60	2.0 × 10 ⁻¹⁶	5.75	0.0165
Biome (B)	0.09	0.7612	0.21	0.6452	0.67	0.4142
D × B	7.03	0.0080	6.51	0.0107	1.26	0.2623
Conditional model	Estin	nate ± se	Estimate ± se		Estimate ± se	
Intercept	1.414	4 ± 0.310	-0.418 ± 0.207 -1.842 ± 0.164 -0.322 ± 0.274 0.642 ± 0.251 Estimate \pm se 1.028 ± 0.140		1.004 ± 0.125 - 0.307 ± 0.124 0.085 ± 0.168	
D (northward)	-1.73	4 ± 0.368				
B (temperate)	0.714	4 ± 0.426				
D×B	-1.31	0 ± 0.494			0.194 ± 0.173	
Dispersion model	Estin	nate ± se			Estimate ± se	
Intercept		-				
D (northward)	-		1.874 ± 0.266		-	
B (temperate)		-	0.75	54 ± 0.198	_	
D×B		-	-1.575 ± 0.389		_	
Random effects	Va	ariance	Variance		Variance	
Plant species: Network).368	3.4 × 10 ⁻⁹		0.157	
Network	C).077		0.174	(0.052

Proportion of plant species interacting with birds during migration (n = 434 observations) (i) (Fig. 2a), frequency of seed-dispersal interactions with birds during migration whenever these interactions occurred (non-zero frequencies; n = 260 observations) out of the total interaction weight (ii) (Fig. 2b) and number of bird species dispersing each plant species during migration whenever these interactions occurred (n = 260 observations) (iii) (Fig. 2c). Family and link functions are shown in parentheses. All models included network identity and plant species nested within network as random factors to account for the repeated measures at these levels. Model (ii) also includes a dispersion model because the dispersion parameter Φ of the beta distribution was allowed to vary in response to the interactive effects of direction and biome⁹⁵. *P* values (two-sided) < 0.05 and significant model estimates (P < 0.05) are shown in bold. In all models, southward and Mediterranean were used as the reference categories (intercepts) for the factors direction (D) and biome (B), respectively.

Extended Data Table 4 | Significance of the fixed factors migration direction and biome, and their interaction, in GLMMs testing effects on the proportion of migratory bird species that were Palaearctic migrants, and in the network-level frequency of seed-dispersal interactions with Palaearctic migrants

Fixed-effects	(<i>i</i>) Proportion of that were P (Binom	 (<i>i</i>) Proportion of migratory bird species that were Palearctic migrants (Binomial, logit link) 		(<i>ii</i>) Interaction frequency during migrations with Palearctic migrants (Beta, logit link)		
Hypothesis testing	X ²	Р	X ²	Р		
Direction (D)	7.98	0.0047	32.47	2.0×10^{-16}		
Biome (B)	9.14	0.0025	12.98	0.0003		
D×B	0.11	0.7458	7.12	0.0076		
Conditional model	Esti	Estimate ± se 0.034 ± 0.363		Estimate ± se 1.188 ± 0.449		
Intercept	0.03					
D (northward)	1.00	1.004 ± 0.466		2.508 ± 0.503		
B (temperate)	1.42	1.429 ± 0.534 0.268 ± 0.825		9 ± 0.531		
D×B	0.26			-1.516 ± 0.568		
Dispersion model	Esti	mate ± se	Esti	mate ± se		
Intercept		-	1.31	8 ± 0.560		
D (northward)		-	3.40	4 ± 0.994		
B (temperate)		-	2.93	6 ± 0.742		
Random effects	V	ariance	V	ariance		
Network		0.336		0.088		

Family and link functions are shown in parentheses. Models included network identity as random factor to account for the repeated measures within networks (n = 26 observations, 13 networks × 2 directions). We used data only from Palaearctic migrants because the frequencies from both migrant types are fully interdependent (Fig. 3a, b). Model (ii) also includes a dispersion model because the dispersion parameter Φ of the beta distribution was allowed to vary in response to the additive effects of direction and biome⁶⁸. *P* values (two-sided) < 0.05 and significant model estimates (P < 0.05) are shown in bold. Results for the species richness of all migrant species pooled are provided in Extended Data Fig. 4.

In all models, southward and Mediterranean were used as the reference categories (intercepts) for the factors direction (D) and biome (B), respectively.

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\boxtimes		For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
	\boxtimes	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
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		Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.

Software and code

Policy information	about <u>availability of computer code</u>
Data collection	Our study includes both data collection (field sampling and molecular analyses) and data compilation from previous studies. We used SEQUENCHER v. 4.9, BioEdit v. 7.0.9 and Chromas v. 2.5.1 for sequence alignment and editing. We used the online platforms BOLD (www.boldsystems.org) and BLAST (https://blast.ncbi.nlm.nih.gov/Blast.cgi) for species identification from amplified DNA sequences (i.e. DNA barcoding).
Data analysis	We conducted all analyses in R version 3.5.2. We used the R packages glmmTMB (v. 0.2.3), car (v. 2.1-6), ape (v. 5.3), phytools (v. 0.6-99), Ime4 (v. 1.1-19) and bipartite (v. 2.13). We made the figures using the R packages 'ggplot2' (v. 3.3.0) and cowplot (v. 0.9.4). The R scripts used to generate all results and figures are available through the Dryad Digital Repository (https://doi.org/10.5061/dryad.15dv41nx3).

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Data

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All data used in the analyses are available through the Dryad Digital Repository (https://doi.org/10.5061/dryad.15dv41nx3). The dated phylogeny of seed plants (Spermatophyta) used to obtain our phylogenetic tree is available through GitHub (https://github.com/FePhyFoFum/big_seed_plant_trees/releases). Data on bird

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Study description	We combined phenological and migration information with data on 949 seed-dispersal interactions between 46 bird and 81 plant species from 13 woodland communities across Europe. The study design is factorial, as we tested the interactive effect between migration direction and community biome (direction x biome) on different response variables. The data structure is hierachical because we have species-level data for plants at birds nested within sites. The data structure also has repeated (paired) measures, with a data points per migration direction (southward and northward).
Research sample	Research sample includes seed-dispersal networks, that is, interacting communities of 46 frugivorous bird species that disperse the seeds of 81 fruiting plant species from 13 woodland communities across Europe (see Extended Data Table 2). Seed dispersal networks are expressed as interaction matrices where each row i represents a plant species and each column j represents a bird species. Elements in the matrices (wij) denote whether pairwise plant-bird interactions were observed (wij > 0) or not (wij = 0) and, if so, their value account for interaction weight. In this case, the quantity of seeds of each plant species dispersed by each bird species.
	The total sample size for plants, the main subject of the study, includes 434 observations (i.e. "plant-species / site / migration" combinations).
	Sex and age of the individuals is not relevant for the purposes of this study.
Sampling strategy	All the study networks (n = 13) were sampled all year-round, a prerequisite to cover the entire fruiting periods of all local fleshy- fruited species, as well as the prenuptial and postnuptial migration periods of all migratory birds. Our study includes a combination of networks from previous studies (n = 5) and newly sampled networks (n = 8). The number of networks was evenly distributed between the Mediterranean (n = 6) and the temperate biomes (n = 7) of Europe, which allowed a sufficient sample size to test for biome effects.
Data collection	Seven of the eight new networks were sampled through field sampling of bird-dispersed seeds and subsequent disperser identification by means of DNA-barcoding analysis. Individual seeds or droppings with seeds were sampled for DNA-barcoding analysis into 1.5- or 2.0-ml sterile tubes that were labelled and stored in a freezer at -20°C until DNA extraction. The molecular analyses for species identification was conducted by J.M.A. and J.C.I.
	The other new network was sampled through dietary analysis of birds captured in mist nets. Captured birds were individually placed in ringing bags for up to 30 min until they ejected droppings.
	The networks compiled from previous studies were obtained either through focal plant observations of birds feeding on fruits or through dietary analysis of birds captured in mist nets. Bird samples were collected in the field by J.P.GV., J.A., J.M.A., R.S.B., T.B., G.EA., N.F., D.G., J.C.I., P.J., P.K., W.J.S., E.V., L.P.d.S. and R.H. The networks from previous studies were compiled by B.I.S. Data on fruiting phenology and bird migrations was gathered by B.R., J.P.GV. and A.T.
Timing and spatial scale	New data in this study was collected between 2012 and 2018. Data available from previous studies were collected between 1981 and 1999. Sampling consisted on periodical surveys during 1–6 years, generally every 2 weeks, in which seed traps were revised, birds were mist-netted or focal plants were observed. Such frequency of sampling during all-year-round allows recording interactions between all bird adn plant species of each community, regardless the season. See details in Table S1. The data was collected in 13 sites across Europe and each site spans several square kilometers.
Data exclusions	No data were excluded.
Reproducibility	The analyses conducted in this study can be reproduced using the R code, which we will made publicly available once the article is accepted for publication. Moreover, we will make available in the DRYAD repository the complete dataset with the detailed information for each sample and its corresponding sequence obtained through DNA barcoding. The sequences include the sample code and all information associated: sampling date, site, seed species and number of seeds of the sample, bird species identified through DNA barcoding, percentage of similarity with best matched sequence and GenBank accession number of best matching sequence. In addition, we keep the DNA aliquots of all DNA extractions from each sample (voucher numbers correspond with 'sample_code') in our labs at University of Oviedo and EBD-CSIC (Spain). These aliquots are available under request. All seed samples are stored by J.P.GV at the laboratory of Botany in the University of Cádiz (Spain).
Randomization	Randomization does not apply as we did not conduct experiments. However, we did assess potential biases regarding the sampling methods of study networks and the approach used to obtain seed-dispersal periods (see Supplementary Discussion 2; Supplementary Figures 5-7).

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Blinding does not apply as we did not conduct expension	riments

Did the study involve field work? X Yes No

Blinding

Field work, collection and transport

Field conditions	We conducted fieldwork on several sites all year round during, for 1-6 years. The field conditions varied seasonally and between sites. The exact field conditions are not relevant because they do not impact the results. In the case of mit-netting, field work was only conducted when it was not raining .
Location	The latitude and longitude of the study sites is provided in Extended Data Table 1.
Access & import/export	 Permits to access to the study sites of the new data collected in this study (see Extended Data Table 1): Garrapilos (Spain): military site; permission given by "Servicio de Cría Caballar de las Fuerzas Armadas" to J.P.GV. (CC-42B00100-S-13-5090; 16 October 2013). Cabañeros (Spain): national park; permission given by the director of the "Cabañeros National Park" to E.V. (June 2016). Arbazal (Spain): private site; permission given by local authority and owners to D.G. (February 2016). Vale Soeiro (Portugal): permission for mist-netting and bird ringing given by ICNF ("Instituto da Conservação da Natureza e das Florestas") to L.P.d.S. (117/2012, 126/2013, 130/2014, 137/2015, 140/2016, 146/2017 and 123/2018). Ficuzza (Italy): private site; permission given by owners to R.S.B. (May 2016). Bradfield Woods (UK): natural reserve; permission given by "Suffolk Wildlife Trust" to J.P.GV. and W.J.S (30 March 2016). Bauerbach (Germany): public site; permission by local authority to N.F. (6 April 2016). Hebdów (Poland): public and private site; permission by local authority and owners to P.K. (May 2016).
Disturbance	Our study did not entail disturbance at the study sites. We placed seed traps in the field sites or used mist-nets, which are placed temporarily and do not cause impacts.

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Ma	terials & experimental systems	Me	thods
n/a	Involved in the study	n/a	Involved in the study
\boxtimes	Antibodies	\boxtimes	ChIP-seq
\boxtimes	Eukaryotic cell lines	\times	Flow cytometry
\boxtimes	Palaeontology and archaeology	\boxtimes	MRI-based neuroimaging
	Animals and other organisms		
\boxtimes	Human research participants		
\boxtimes	Clinical data		
\boxtimes	Dual use research of concern		

Animals and other organisms

Policy information about studies involving animals; ARRIVE guidelines recommended for reporting animal research

Laboratory animals	No laboratory animals were used in the study.
Wild animals	For sampling the seed-dispersal network from Portugal (Vale Soeiro), birds were caught in the field using mist-nets and immediately released in the same location after they ejected droppings in the cloth bag. No bird was injured, killed or kept captive.
	Overall, 45 species were captured in a total of 4462 bird captures including 3123 individuals. In 1330 captures (846 individuals) of such captures, the bird produced droppings with seeds (13 bird species).
	Full list of bird species captured (n = 45 species): Accipiter nisus, Acrocephalus scirpaceus, Aegithalos caudatus, Alcedo atthis, Anthus trivialis, Caprimulgus europaeus, Caprimulgus ruficollis, Carduelis chloris, Carduelis spinus, Certhia brachydactyla, Cyanistes caeruleus, Dendrocopos major, Erithacus rubecula, Ficedula hypoleuca, Fringilla coelebs, Fringilla montifringilla, Garrulus glandarius, Hippolais polyglotta, Lophophanes cristatus, Luscinia megarhynchos, Muscicapa striata, Parus major, Passer domesticus, Periparus ater, Phoenicurus ochruros, Phoenicurus phoenicurus, Phylloscopus collybita, Phylloscopus ibericus, Phylloscopus trochilus, Picus sharpei, Prunella modularis, Pyrrhula pyrrhula, Regulus ignicapillus, Serinus serinus, Streptopelia turtur, Sylvia atricapilla, Sylvia borin, Sylvia cantillans, Sylvia communis, Sylvia melanocephala, Sylvia undata, Troglodytes troglodytes, Turdus iliacus, Turdus merula and Turdus philomelos.
	List of bird species that produced droppings with seeds (n = 13 species): Cyanistes caeruleus, Erithacus rubecula, Ficedula hypoleuca, Parus major, Phoenicurus phoenicurus, Sylvia atricapilla, Sylvia borin, Sylvia cantillans, Sylvia communis, Sylvia melanocephala, Sylvia undata, Turdus merula and Turdus philomelos.

Sex and age of the individuals is unknown (and not relevant for this study).

Field-collected samples Seed samples collected in the field were stored in the freezer at -20°C until DNA-barcoding analyses.

Ethics oversight No ethical approval was required as no bird was killed, injured or kept captive and we used normal procedures for mist-netting.

Note that full information on the approval of the study protocol must also be provided in the manuscript.