EuPPollNet: A European database of plant-pollinator networks

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Abstract

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Motivation: Pollinators play a crucial role in maintaining Earth's terrestrial biodiversity and human food production by mediating sexual reproduction for most flowering plants. Indeed, the network of interactions formed by plants and pollinators constitutes the backbone 89 of plant-pollinator community stability and functioning. However, rapid human-induced envi-90 ronmental changes are compromising the long-term persistence of plant-pollinator interaction 91 networks. One of the major challenges for pollinator conservation is the lack of robust gen-92 eralisable data capturing how plant-pollinator communities are structured across space and 93 time. Here, we present the EuPPollNet (European Plant-Pollinator Networks) database, a fully open and reproducible European-level database containing harmonized taxonomic data 95 on plant-pollinator interactions referenced in both space and time, along with other ecolog-96 ical variables of interest. This database offers an open workflow that allows researchers to 97 track data-curation decisions and edit them according to their preferences. We present the 98 taxonomic and sampling coverage of EuPPollNet, and summarize key structural properties gq in plant-pollinator networks. We hope EuPPollNet will stimulate future research that fills 100 the taxonomic, ecological, and geographical data gaps on plant-pollinator interactions that we 101 have identified. Further, the variation in the structure of the networks in EuPPollNet pro-102 vides a strong basis for future studies aimed at quantifying drivers of plant-pollinator network 103 change and guiding future conservation planning for plants and pollinators. 104

Main Types of Variables Included: EuPPollNet contains 1,174,535 interactions between plants and pollinators from 1,876 distinct networks (i.e., distinct sampling event in space or time), which belong to 54 different studies distributed across 23 European countries. In addition, information about sampling methodology, habitat type, bio-climatic region, and further taxonomic rank information for both plant and pollinator species are also provided (i.e., genus, family and order).

Spatial location and grain: The database contains 1,214 different sampling locations from natural and anthropogenic habitats that fall in 8 different bio-climatic regions. All records are geo-referenced and presented in the World Geodetic System 1984 (WGS84).

Time period and grain: Species interaction data was collected between 2004 and 2021. All records are time-referenced and most of the studies documented interactions within a single flowering season (68.52%).

Major taxa and level of measurement: The database contains interaction data at the species level for 94.22% of the records, including a total of 1,411 plant and 2,223 pollinator species. The database covers 5.56% of the European species of flowering plants, 34.38% of bees, 26.21% of butterflies, and 33.63% of syrphid species at the European level.

Software format: The database was built with the R programming language and is stored as ".rds" and ".csv" formats. The construction of the database is fully reproducible and can be accessed at the following link: https://github.com/JoseBSL/EuPPollNet.git.

124 KEYWORDS

- Angiosperms, connectance, flowering plants, nestedness, plant-pollinator networks, pollinators, species
- 126 interactions

1 | INTRODUCTION

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Plant-pollinator interactions involve a great diversity of species, largely attributed to their co-128 evolutionary history (Ollerton, 2017), and are critically important for terrestrial biodiversity 129 and economic productivity. The synergistic effects of climate change with other global change 130 pressures are threatening worldwide biodiversity (Bellard et al., 2014; Sala et al., 2000), includ-131 ing plant and pollinator species as well as their interactions (Eichenberg et al., 2021; Goulson, 132 Nicholls, Botías, & Rotheray, 2015; Settele, Bishop, & Potts, 2016). Under this scenario, the 133 increasing availability of biodiversity data plays a major role in our ecological understand-134 ing of species status, trends, and conservation (Heberling, Miller, Noesgaard, Weingart, & 135 Schigel, 2021; Zattara & Aizen, 2021). However, our knowledge of plant and pollinator species 136 and their network of interactions still exhibits major temporal, spatial and taxonomic biases 137 (Archer, Pirk, Carvalheiro, & Nicolson, 2014; Marshall et al., 2024; Poisot et al., 2021; Troia 138 & McManamay, 2016), limiting our ability to effectively protect their biodiversity. 139

The interactions between different plant and pollinator species within a community form complex networks. Macro-ecological analyses of the topology of these networks have revealed common properties, such as truncated power-law degree distributions (Jordano, Bascompte, & Olesen, 2003) or nestedness (Bascompte, Jordano, Melián, & Olesen, 2003). Large-scale analyses across multiple studies can quantify patterns across geographic regions (Olesen & Jordano, 2002; Traveset et al., 2016) or environmental gradients (Ramos-Jiliberto et al., 2010; Rech et al., 2016; Saunders et al., 2023) that cannot be examined in a single study. Although macro-ecological approaches that use ecological interactions make significant contributions to knowledge (Windsor, Hoogen, Crowther, & Evans, 2023), such synthesis work must consider variation across studies in the spatio-temporal nature of the data (Burkle & Alarcón, 2011; García et al., 2024). For instance, plant-pollinator studies tend to differ in sampling effort and methodology which affect the structure of the resulting plant-pollinator networks (Gibson, Knott, Eberlein, & Memmott, 2011; Jordano, 2016; Schwarz et al., 2020). Most plantpollinator networks have unobserved interactions (Chacoff et al., 2012; Olesen et al., 2011), and thus research that attempts to synthesize across published studies must have access to raw data on interactions in order to statistically account for sampling effort and completeness. This emphasizes the importance of providing data in its rawest possible form in datasets that will be utilized for synthesis and macro-ecological studies.

Europe is one of the continents with a larger amount of available biodiversity data (Proença et al., 2017), yet still exhibits major gaps (Bennett et al., 2018; Wetzel et al., 2018). While species checklists need to be treated carefully, especially at a macro-ecological scale (Grenié et al., 2023), the growing number of European plant and pollinator checklists (Reverté et al., 2023), along with occurrence data (Zattara & Aizen, 2021), is setting a foundation for the conservation of its flora and their pollinators. However, species richness is just one component of biodiversity and documenting the interaction between plants and pollinators is essential for understanding biodiversity change (Jordano, 2016). Numerous works have studied plant-pollinator interactions in the last decades, generating thousands of plant-pollinator

interaction networks worldwide. Several initiatives have tried to integrate plant-pollinator interaction data into databases such as Mangal (Poisot et al., 2016) or GloBI (Poelen, Simons, & Mungall, 2014), resulting in numerous large scale comparative analyses that have enhanced our understanding of the ecology of plants and pollinators (e.g., European wild bee data trends; Marshall et al., 2024). Despite all these resources, Europe lacks accessible harmonized plant-pollinator interaction data that allow researchers to evaluate plant-pollinator interactions at a European level. For example, only over a dozen of European plant-pollinator networks are included in Mangal, while GloBI focuses on pairwise interactions disconnected from the com-munity context. Assembling and curating the existing information on EU plant-pollinator networks will guide research efforts, conservation planning and will set a foundation for future global change research.

Here, we present the European Plant-Pollinator Networks database (EuPPollNet), which contains harmonized information on plant-pollinator interactions at the European level. The pollinator taxonomic groups include the main orders of entomofauna that visit and pollinate flowering plants in Europe. These comprise insect species from the orders Hymenoptera, Diptera, Lepidoptera, and Coleoptera, accounting for almost the totality of recorded interactions in EuPPollNet (99.84%). To understand the scope of the database, we examined the taxonomic and sampling coverage of the different plant and pollinator species at the European level with the help of the most up-to-date species checklists and rarefaction analyses. In addition, for bees and plants, we evaluated if there is a phylogenetic signal in the presence-absence of interaction data.

EuPPollNet contains one of the largest sets of plant-pollinator networks collated to date, providing a unique opportunity to examine the prevalence of key structural metrics across networks. For example, despite the large theoretical literature on the meaning of a nested structure in plant-pollinator networks (Bascompte & Jordano, 2007; Guimaraes, 2020), where generalist species interact with all other species but specialists interact preferentially only with generalists, this pattern has only been empirically evaluated with a relatively small number of networks (Bascompte et al., 2003; Payrató-Borras, Hernández, & Moreno, 2019; Staniczenko, Kopp, & Allesina, 2013), and is still debated how structural metrics such as connectance and nestedness change across latitudes and bioclimatic regions (Olesen & Jordano, 2002; Song, Rohr, & Saavedra, 2017; Trøjelsgaard & Olesen, 2013).

Overall, EuPPollNet aims to cover a wide range of taxonomic groups and habitats, while also providing other variables of interest that define the ecological context and sampling methods of the study. In addition, EuPPollNet offers a transparent and accessible workflow of its data management and species harmonization that allows the database to be reused and to expand over time. This database provides a large number of community-level networks with curated and harmonized data, distinguishing it from other currently available resources that contain plant-pollinator interactions. We expect that EuPPollNet can be used to evaluate macro-ecological processes in plant-pollinator networks, guide conservation planning, and set a baseline for global change research.

2 | METHODS

Data acquisition

The EuPPollNet database includes published and unpublished studies compiled initially by a 209 wide number of researchers and institutions within the European continent as defined by the European Environment Agency (Stanners & Bourdeau, 1995). As this database is the result 211 of one of the work packages of the European project Safeguard (Safeguarding European wild 212 pollinators; https://doi.org/10.3030/101003476), first, data was directly requested from mem-213 bers of the Safeguard project in May 2022. Second, the request was extended to data owners 214 outside of the project, with data collection concluding in August 2024. These other data own-215 ers were identified by direct communication with colleagues suggested by Safeguard members 216 and by directly searching for studies on Google Scholar of under-represented regions within the 217 database. While Google Scholar lacks reproducibility (Gusenbauer & Haddaway, 2020), it still 218 remains the most comprehensive search engine to date (Gusenbauer, 2019). This approach 219 maximized the potential number of studies that could be incorporated in this database. The 220 search strings used were "plant-pollinator interactions" and "plant-pollinator networks". To 221 maintain high quality standards that will support robust future ecological research, we only 222 included studies that met the following criteria: 1) studies containing time- and geo-referenced 223 records of plant-pollinator interactions; and 2) studies that quantify interactions by document-224 ing them as the contact between a pollinator and the reproductive structure of a specific plant being sampled (i.e., phyto-centric networks). 226

227 Dataset description

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The database contains 54 independent published and unpublished studies conducted during the time period 2004 - 2021 in 23 different countries (Figure 1a and Figure 1b). studies differ in sampling effort and methodology, and thus documenting sampling methods and sampling effort is an important feature of EuPPollNet. Most studies took place within a single flowering season (68.52%), sampled a given location for an average of 7.11 days, and documented only diurnal interactions, with transects being the most common sampling method (64.81%). All the studies documented interactions with Hymenopterans (with 50.00%) considering all Hymenopterans, 46.15% only wild bees and 3.85% only bumblebees), 92.31% documented interactions with Dipterans (with 46.15% considering all Dipterans, 46.15% only syrphids and 5.77% recorded syrphids plus bombylids or tachinid flies), 63.46% with Lepidopterans, and 32.69% with Coleopterans. The database includes a total of 1,174,535 distinct interactions. Most of the pollinator species belong to the orders Hymenoptera, Diptera, Lepidoptera and Coleoptera (89.11%), which account for nearly all interactions in this database (99.84%). Species that belong to other orders 10.89% are not explored in this study as they conduct a minor fraction of the total interactions (0.16%). Hymenoptera and Diptera contain the highest number of species comprising each approximately 1,000 species in the database. However, the majority of plant-pollinator interactions are from Hymenoptera species (89.59%; Figure 1c). Notably, the western honey bee, *Apis mellifera*, represents 69.36% of the total interaction records from the database and an average of 30.33% of the total interactions per network.

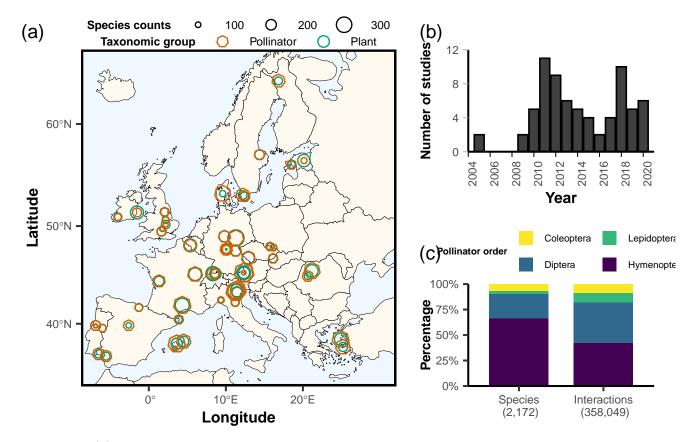


Figure 1. (a) Locations of the studies in EuPPollNet across the European continent, showing the total number of pollinator (i.e., orange heptagon) and plant (i.e., green circles) species per study. The sizes of these shapes are proportional to the respective species counts. For visualization purposes, we have selected only a single location per study. (b) Number of studies by year in EuPPollNet. (c) Proportion of species and interactions across the four main pollinator orders in EuPPollNet, excluding interactions from *Apis mellifera*. The total number of species and interactions is indicated in parentheses at the bottom.

48 Data structure

The EuPPollNet database is available in both .csv and .rds formats and contains a total of 31 columns (Table 1), where each row represents a single interaction between a plant and a 250 pollinator species. These columns include information about the study and network identifiers 251 (columns 1 and 2), sampling method (3), habitat type as described by the author, and a 252 unified habitat classification across studies (4 and 5), bioregion where the network is located 253 (6), country, locality, and latitude-longitude coordinates (7 to 10), date of the interaction 254 (11), number of interactions (12), taxonomic information about plants (13 to 20), taxonomic 255 information about pollinators (21 to 28), and information about the availability of floral count 256 data (29). The flower count data is provided in a separate file (.csv or .rds) and can be 257 merged with the interaction data through the "Flower data merger" column (30). Note 258 that although two-thirds of studies include information on floral abundance, the methods and 259 units vary greatly across studies. To construct a plant-pollinator network matrix within a 260 single flowering season at the site level, users should group interactions by plant and pollinator 261 species, site, study, and year. Finally, metadata at the study level is provided in a separate 262 file, including information about the authors, digital object identifier (if available), sampling 263 time, and taxonomic coverage of the main pollinator groups for each study. 264

Table 1. Column names and their descriptions within the EuPPollNet database.

Variable	Description
Study_id Network_id Sampling_method Authors_habitat EuPPollNet_habitat	Identifier of the study Identifier of a site sampled within a study Type of plant-pollinator sampling Type of habitat as described by the authors Type of habitat homogenized across studies
Bioregion Country Locality Latitude Longitude	European biogiographical regions Country where the plant-pollinator interaction was observed Locality where the plant-pollinator interaction was observed North-south position of the observed interaction in decimal degrees East-west position of the observed interaction in decimal degrees
Date Interaction Plant_original_name Plant_accepted_name Plant_rank	Year, month and day when the observation took place Number of interactions. By default is 1 as interactions are provided ungrouped Plant species name given by the authors Harmonized plant species name in the database Taxonomic rank of the observation
Plant_order Plant_family Plant_genus Plant_unsure_id Plant_uncertainty_type	Order taxonomic rank of the observed plant species Family taxonomic rank of the observed plant species Genus taxonomic rank of the observed plant species Category to indicate if the plant species name is unsure (Yes) or not (No) If the name is unsure, type of species uncertainty is provided
Pollinator_original_name Pollinator_accepted_name Pollinator_rank Pollinator_order Pollinator_family	Pollinator species name given by the authors Harmonized pollinator species name in the database Taxonomic rank of the observation Order taxonomic rank of the observed pollinator species Family taxonomic rank of the observed pollinator species
Pollinator_genus Pollinator_unsure_id Pollinator_uncertainty_type Flower_data Flower_data_merger	Genus taxonomic rank of the observed pollinator species Category to indicate if the pollinator species name is unsure (Yes) or not (No) If the name is unsure, type of species uncertainty is provided Floral data availability (Yes) or (No) Column to merge floral data with the interaction dataset

Taxonomic harmonization

All plant and pollinator species names were checked and harmonized in R using **rgbif** (Chamberlain, Oldoni, & Waller, 2022). The protocol for plants and pollinators is similar but slightly different given the availability of the different taxonomic resources. For transparency, we have included in the database the original species name, the new assigned name, and, if the name of the species is uncertain (e.g., species complex or species alike). In addition, taxonomic information at genus, family and order level was downloaded for each species.

For plants: (i) we initially verified the exact matches against the GBIF species checklist; (ii) we selected unmatched cases and fixed orthographic errors; (iii) we retrieved again taxonomic

information for those unmatched cases, evaluated accuracy of fuzzy matching and programmatically fixed records that are still not found; (iv) finally, we used the World Flora Taxonomic
Backbone (Govaerts, Nic Lughadha, Black, Turner, & Paton, 2021; WFO, July 7, 2022) as
the ultimate filter for taxonomic information as we used it to calculate the plant taxonomic
coverage of our database.

For pollinators: (i) we first created a checklist of species names for the most representative 279 pollinator groups at the European level by combining the most up to date published checklists 280 of bees and syrphids (Reverté et al., 2023), and butterflies (Wiemers et al., 2018); (ii) we 281 compared pollinator species names against the checklist and recovered some unmatched cases 282 with restrictive fuzzy matching by using **stringdist** package (Van der Loo et al., 2014); (iii) 283 we fixed unmatched records when necessary and retrieved the taxonomic information for all 284 species from GBIF; (iv) we fixed the non-found cases in the GBIF checklist and made sure that 285 all species names from bees, syrphids and butterflies were named according to their respective 286 species checklists. Coleoptera species names were only checked against the GBIF checklist.

288 Taxonomic coverage

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To assess the completeness of plant and pollinator species in the EuPPollNet database at the European level, we used the aforementioned checklists for plants and pollinators. Specifically for plants, we refined the checklist to include only plants occurring in Europe and excluded taxonomic groups not associated with biotic pollination. We did this by first excluding the families considered to have exclusively a wind pollination mode (see Culley, Weller, & Sakai, 2002), and then by filtering out the genera with wind or non-biotic pollination from families that exhibit both biotic and non-biotic pollination modes. Additionally, we manually included exotic species and added unresolved species names that were not present in the accepted names of the checklist at the current version of usage. For pollinators, we compared only the taxonomic coverage of bees, syrphids and butterflies at the European level, as these are the only taxonomic groups with species checklists available in Europe. In addition, the potential number of pollinator species in Europe was estimated by combining the total number of species of bees, syrphids, butterflies and species belonging to other taxonomic groups. The total number of species from other taxonomic groups was extrapolated by assuming that their coverage within the database is equal to the mean coverage of bees, syrphids and butterflies (mean coverage = 31.4; sd = 4.51).

Finally, to evaluate if the presence-absence of interaction records for bees and flowering plants follows a phylogenetic pattern within the database, we calculated its phylogenetic signal at genus and family level, respectively. The phylogenetic signal was calculated by using the phylosig function from the **phytools** package (Revell, 2012). We extracted the phylogenetic information for bees from a genus level phylogeny (Hedtke, Patiny, & Danforth, 2013) and processed it using the packages ape (Paradis et al., 2019), MCMCglmm (Hadfield, 2010) and **phytools**. For plants, the phylogenetic tree was obtained from a species level plant phylogeny (Smith & Brown, 2018) with the help of the **rtree** package (Li, 2023).

313 Sampling coverage

The completeness of the EuPPollNet database was evaluated by exploring the rarefied accumulation curves of plant and pollinator species and their interactions across the different networks.

In addition, we computed the accumulation curve of pollinator species with an increasing number of plant species as an indicator for how many pollinator species are likely responsible for the pollination of flowering plants (e.g., Kleijn et al., 2015 for crops). The rarefied and extrapolated sampling curves were obtained using the iNEXT package (Hsieh & Chao, 2016). The different rarefied curves were complemented with 100 bootstrapped accumulation curves.

321 Habitat type and bioclimatic region

We describe the habitat type for each site using information from Corine Land Cover (CLC, 322 version 2018) extracted using the Terra package (Hijmans et al., 2022), visual inspection of 323 Google Earth imagery and the habitat classification from the authors. These different habitat 324 categories (see definitions in supplementary text) allow a quick comparison and understanding 325 of the habitat types from the database. Moreover, Europe is characterized by a great variety 326 of environmental conditions that harbor different biota. Thus, to allow authors to explore 327 the set of studies that share similar environmental conditions and species, we assigned a 328 biogeographical region to each site. The biogeographical regions were downloaded from the 329 European Environment Agency (version 2016) and were matched to the different sites using a 330 spatial joint from the sf package (Pebesma et al., 2018). 331

332 Network analyses

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To provide a general overview of the structure of plant-pollinator networks in EuPPollNet, we quantified connectance and nestedness for each network, and examined how these network metrics change across latitude of studies and bioclimatic regions in Europe. We selected these two network metrics as they are commonly evaluated in plant-pollinator network studies and capture structural properties with a straightforward interpretation. We implemented "standardised" versions of connectance and nestedness to account for the effect of sampling effort on network metrics. As connectance is negatively associated with network size (Jordano, 1987), we evaluated how network connectance was associated with the number of species (i.e., log of geometric mean of plants and pollinators) and extracted the residuals from this association (i.e., residual connectance) as a measurement of corrected connectance. The relationship between residual connectance and species richness was investigated using a beta-regression implemented with the package **betareg** (Cribari-Neto & Zeileis, 2010). We used NODFc to compare nestedness across networks, as it corrects by connectance and the number of species (Song et al., 2017). This metric was calculated using the maxnodf package (Hoeppke & Simmons, 2021). Both residual connectance and NODFc were used as dependent variables to evaluate their association with latitude. In addition, we quantified the association between

connectance and nestedness with the number of species per network using a Kendall rank correlation coefficient to compare the strength of associations between network structures and species richness across both network metrics.

Finally, to compare if networks are more or less nested than expected by chance, we employed 352 the traditional z-score approach with the widely used nestedness metric (NODF) from Almeida-353 Neto, Guimaraes, Guimaraes Jr, Loyola, & Ulrich (2008). The z-score approach allows us to 354 compare our results with previous published nestedness analyses in plant-pollinator networks 355 and only compares each unique network against their randomized versions. We calculated 100 356 null models for each network with the help of the vaznull function from the package bipartite 357 (Dormann, Gruber, & Fründ, 2008). These null networks have the same connectance and 358 number of plant and pollinator species as the empirical ones, but different marginal totals. 359 Both connectance and nestedness (NODF) were estimated for each network by using the 360 function *networklevel* from **bipartite**. 361

3 | RESULTS

Taxonomic coverage

Europe hosts approximately 5,000 species of pollinators and 25,000 species of plants that 364 benefit from animal pollination. EuPPollNet contains a total of 2,223 pollinators and 1,411 365 plant species. The coverage of the main pollinator groups occurring in Europe is 34.38% for 366 bees, 33.63% for syrphids and 26.21% for butterflies (see Figure S1 for coverage at the family 367 level for bees and butterflies, and at the subfamily level for syrphids). Bees constitute 89.05%368 of the interactions in EuPPollNet, and 77.89% of the interactions when excluding honey bees. 369 Within the database, 83.82% of bee genera have at least one species with interaction records, 370 and the average coverage of species at the bee genus level is 36.99% (Figure 2). The presence 371 or absence of interaction records for bees does not follow a phylogenetic pattern ($\lambda = 0.07$; 372 P = 0.65). The database coverage of all flowering plant species occurring in Europe is 5.56% 373 (Figure 3), with an average coverage of 9.04% at the plant family level. Approximately, half of 374 the plant families have at least one species with interaction records (52.56%), and the presence 375 or absence of interaction data for the different plant species also does not follow a statistically 376 relevant phylogenetic pattern ($\lambda = 0.26$; P = 0.07). 377

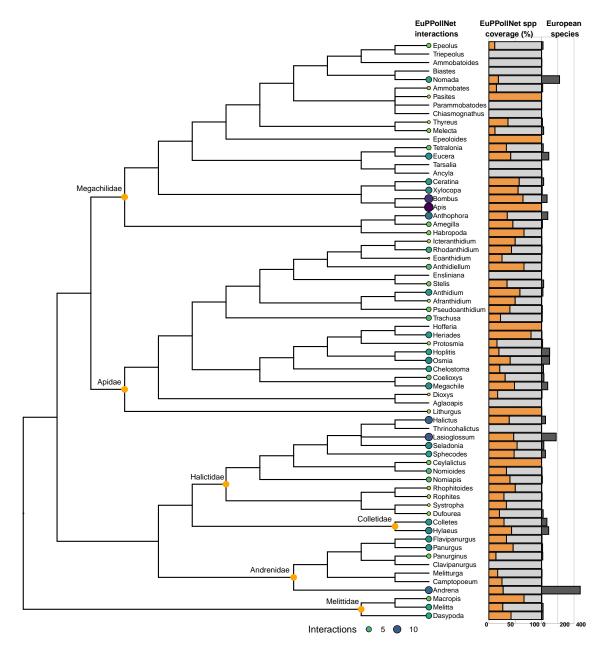


Figure 2. Phylogenetic and taxonomic coverage of the bee genera at European level. The number of interactions recorded per genus in the database is illustrated using circles, with their sizes proportional to the number of interactions on a logarithmic scale. Additionally, a gradient of colors ranging from yellow to dark purple aids in this visualization. The coverage of species recorded in EuPPollNet per genus at the European level is depicted with orange and light grey bars, representing the percentage of species included and not included in the database, respectively, out of the total number of bee species per genus at the European level. Dark grey bars indicate the total number of species per genus at the European level.



Figure 3. Phylogenetic and taxonomic coverage of the plant families at European level. The number of interactions recorded per genus in the database is illustrated using circles, with their sizes proportional to the number of interactions on a logarithmic scale. Additionally, a gradient of colors ranging from yellow to dark purple aids in this visualization. The coverage of species recorded in EuPPollNet per genus at the European level is depicted with orange and light grey bars, representing the percentage of species included and not included in the database, respectively, out of the total number of flowering plant species per family at the European level. Grey bars indicate the total number of species per family at the European level on logarithmic scale.

378 Sampling coverage

The estimated sampling coverage of plant and pollinator species within EuPPollNet across 379 the different networks is approximately 97% for both taxonomic groups. This suggests that 380 the rarefied accumulation curves of both plant and pollinator species exhibit already a "quasi-381 asymptotic" growth of species richness by considering the current number of networks (Figure 382 4a-4b). The predicted observed species richness by doubling the sampling effort on the already 383 sampled habitat types within the database will only increase pollinator richness by 24.13% and 384 plant richness by 21.14%. However, the sampling coverage of interactions is 74.17%, and by 385 doubling the sampling effort the predicted number of unique interactions recorded will have 386 approximately a twofold increase (54.83%; **Figure 4c**). When we consider the accumulated 387 pollinator richness across sampled plant species, this curve also shows a "quasi-asymptotic" 388 growth with a sampling coverage value of 96.66%. The predicted recorded pollinator species 389 by doubling the number of plants sampled is expected to increase by 21.90% (Figure 4d). We 390 find that a small portion of plant species and pollinator species are shared across a broad range 391 of networks and that most plant (85.05%) and pollinator (86.59%) species are exclusively found 392 in less than 1% of networks (Figures 4e-4f). The most common plant (Trifolium pratense) 393 and pollinator (Bombus pascuorum when excluding Apis mellifera) species are found in 35.69% 394 and 62.18% of networks, respectively.

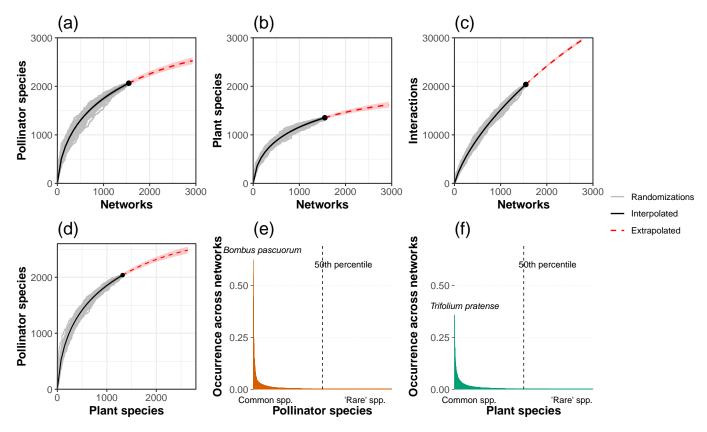


Figure 4. Graphs (a-b-c) indicate the accumulation curves for pollinators, plants, and their interactions across networks. Grey solid lines represent 100 randomized accumulation curves, the black solid lines represent the interpolated curve (i.e., the mean across curves), and the red dashed lines illustrate the extrapolated curve for approximately 3000 networks. The solid black points indicate the number of species and interactions contained in the database. Graph (d) shows the accumulation curve of pollinator species across an increasing number of plant species. This last graph uses the same color and shape structure as the ones in the top panel. Graphs (e-f) indicate the percentage of occurrence (i.e., incidence) of plant and pollinator species across networks. Species on the left (i.e., common) are found in many networks, while species on the right (i.e., rare) are found in few or only a single network. Note that indeed Apis mellifera is the most common pollinator but was excluded from this visualization.

Habitat type and bioclimatic region

The proportion of species from the major pollinator orders within the database differed across 397 habitats and bioclimatic regions (Figure 5). As expected, Hymenoptera was the main taxo-398 nomic order on the majority of habitats, exceeded only by Diptera on the habitat categories 399 of riparian vegetation, moors and heathland, and alpine grasslands. Overall, the proportion of 400 Lepidoptera and Coleoptera species was low across all habitats but Coleoptera were notably 401 more abundant in sclerophyllous vegetation and beaches, dunes and sands habitat categories. 402 Similar patterns were observed when exploring the pollinator proportions by bioclimatic region. 403 In this case, Hymenoptera was the predominant order in all bioclimatic regions, with Diptera 404 being particularly abundant in the Alpine and Atlantic regions. Lepidoptera showed low pro-405 portions across all bioclimatic regions and Coleoptera were only relevant in the Mediterranean 406 region at European level. Notably, the number of studies (Figure 5) and sampling sites 407 (Figure S2) also differed across habitats and bioclimatic regions. The habitats sampled by 408 a higher number of studies in the database were intensive grasslands (28), semi-natural grass-409 lands (12) and sclerophyllous vegetation (10). However, the habitats that contain a higher 410 number of sampling sites were intensive grasslands (620), agricultural margins (432) and agri-411 cultural land (141). The bioclimatic regions with a higher number of studies were Continental 412 (23), Atlantic (13) and Mediterranean (13); and those that contain a higher number of sampling 413 sites were Continental (482), Atlantic (459) and Boreal (439). 414

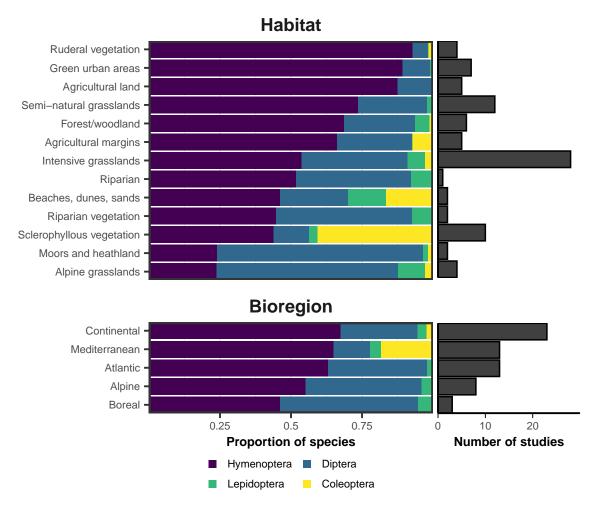


Figure 5. Proportion of species from the major pollinator orders by habitat types and bioclimatic regions in the EuPPollNet database. The orders, from left to right, include Hymenoptera, Diptera, Lepidoptera and Coleoptera. The horizontal barplot on the right indicates the number of studies that were conducted on each habitat type or bioclimatic region. Note that a single study can contribute to more than one habitat or bioclimatic region. Areas with a greater number of studies are more likely to depict accurate proportions of the different pollinator orders in those systems.

Network properties

Connectance values ranged between 0.03 to 0.4 ($\bar{x} = 0.14$) and as expected, followed a negative exponential relationship with the number of species per network (Kendall $\tau = -0.75$, P < 0.01; 417 Figure 6a). Nestedness values (NODFc) ranged between 1.34 to 7.94 ($\bar{x} = 2.81$), and as 418 expected for this metric, were independent of the mean number of species (Kendall $\tau = -$ 419 0.06, P = 0.03; Figure S3). We found that only 31.62% of networks were statistically more 420 nested than expected by chance, with 68.38% showing no statistical difference, and none being 421 less nested than null expectations (Figure 6b). Latitude explained little of the observed 422 variability of residual connectance and nestedness across networks (connectance: $R^2 = 0.02$, P 423 = 0, Figure 6a; NODFc: $R^2 = 0.01$, P = 0.02, Figure 6b). Overall, networks towards higher 424 latitudes showed lower residual connectance and higher nestedness than networks located at 425 lower latitudes. Note that residual connectance and normalised nestedness showed a moderate 426 negative correlation (Kendall $\tau = -0.46$, P = 0). 427

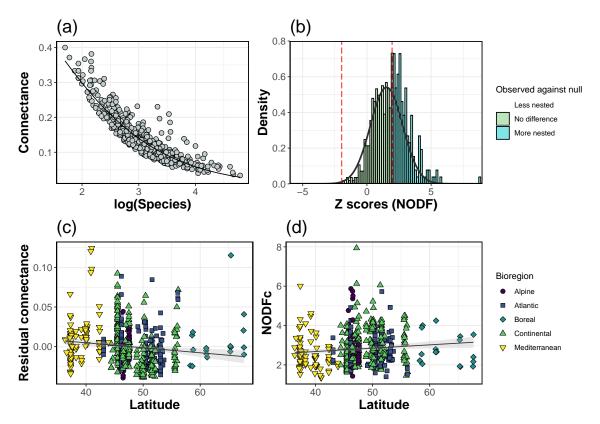


Figure 6. Graph (a) shows the association between network connectance and the geometric mean of plant and pollinator species per network on a log-scale with the respective fitted line from a Beta regression. Graph (b) shows the distribution of z-scores when comparing the nestedness from the empirical networks with their randomised counterparts (100 null models for each network). The vertical red dashed lines represent the z critical value for a two tailed test with alpha = 0.05. Z-scores to the left of the first vertical red dashed line indicate that networks are less nested than expected by chance (red), those between the two dashed lines indicate no statistical difference from random expectations (green), and those to the right indicate that networks are more nested than expected by chance (blue). Graphs (c-d) show the fitted regression of residual connectance and nestedness across the latitudinal range of the studies from the database. The bioclimatic region of each network is indicated with points of different shapes and colours.

28 3 | DISCUSSION

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EuPPollNet offers the largest set of plant-pollinator studies and networks compiled to date 429 at European level. The database contains 1,411 plant and 2,223 pollinator species with over 430 a million interaction records. While the overall sampling coverage of species and interactions 431 is relatively high across the sampled sites, the taxonomic coverage of plants and the main 432 pollinator groups at the European level is still relatively low (i.e., 5.6% for flowering plants 433 and 34.4% for bee species). This likely reflects the fact that most plant and pollinator species 434 are rare and geographically restricted. For example given that most of the plant-pollinator 435 networks from the database are sampled on intensive grasslands, and habitat heterogeneity is a 436 crucial factor in understanding pollinator diversity at European level (Hass et al., 2018; Kleijn 437 et al., 2015; Martínez-Núñez et al., 2022), adding studies on other habitat types is likely to 438 result in a rapid increase of the coverage of plant and pollinator species and their interactions. 439 Indeed, plant and pollinator species were rarely shared across multiple sites, indicating that 440 there are few "common" species and many "rare" ones at the metaweb or continental level. 441 This high number of "rare" species results in an upward slope of the species or interaction 442 accumulation curves (Thompson & Withers, 2003). In other words, minimal sampling efforts 443 are capturing a substantial number of species and interactions, but achieving a comprehensive 444 inventory will require numerous sampling events within and across habitats, particularly for 445 plant-pollinator interactions. 446

Bees are responsible for the majority of the interactions at the metaweb level, but their relevance for network topology changed across habitats and bioclimatic regions. For instance, plant-pollinator communities in the Mediterranean were dominated by bees, while communities in Alpine or Boreal regions were fly species rich or fly-dominated. These patterns are consistent with our current understanding of bee diversity, which peaks in dry or temperate areas (Leclercq et al., 2023; Orr et al., 2021); and with the fact that colder environments (i.e., altitude and latitude wise) harbor a larger fraction of fly pollinators compared to other taxa (Elberling & Olesen, 1999; Lefebvre, Villemant, Fontaine, & Daugeron, 2018). In addition, beetles were only common floral visitors in the Mediterranean region. Although the networks from the database are visitation networks and do not capture pollinator efficiency (Ballantyne, Baldock, & Willmer, 2015), the high proportion of beetles as floral visitors in the Mediterranean provides further support for their potential role as pollinators in this region (Herrera, 2019; León-Osper & Narbona, 2022). The number of butterfly species and interactions were relatively low compared to the other taxa. While Europe contains fewer butterfly species than other regions of the world (Ollerton, 2017), their relevance as pollinators is likely underestimated within this database. This is because a large fraction of studies ($\sim 40\%$) did not sample butterflies, and conventional sampling methods for monitoring other insect pollinators (e.g., bees or flies) may be inadequate for sampling plant-butterfly interactions (Isaac et al.. 2011). Honey bees were present in 87% of networks and conducted on average a third of the total interactions per network. The proportion of honey bees in networks across Europe is higher than in natural communities across the world (~13%; Hung, Kingston, Albrecht, Holway, & Kohn, 2018), potentially reflecting the dominance of intensive grasslands habitats in

EuPPollNet, the highly generalised nature of honeybees, the important role that honeybees are playing as pollinators in Europe, and/or because beekeeping is widely practiced in Europe (Herrera, 2020; Magrach, González-Varo, Boiffier, Vilà, & Bartomeus, 2017; Steffan-Dewenter & Tscharntke, 2000).

Although Europe contains a much larger number of flowering plants than pollinator species (~5 to 1 ratio according to our extrapolation from checklists), the observed number of pollinator species in the database was almost double that of the plants. This could be explained by the fact that all networks are phytocentric, resulting in sampling bias towards pollinator species (Jordano, 2016; Vizentin-Bugoni et al., 2018). While animal-centered sampling is likely to increase the plant-pollinator species ratio (e.g., Encinas-Viso et al., 2023), the spatial scale and environmental context of the sampled communities will also influence their observed diversity, especially given the ability to move of pollinators and the sessile nature of plants. In addition, we found that the accumulation curve of pollinators per plant species does not saturate, which indicates low redundancy of pollinators and that many are regionally "rare". Rare pollinators are functionally important for plant species at large scales (Simpson et al., 2022; Winfree et al., 2018), highlighting the need to conduct further sampling events to observe these rare species and to effectively understand and protect plant-pollinator biodiversity.

Consistent with Olesen & Jordano (2002), we find that residual connectance (i.e., the deviation from the expected connectance for a given network size) was lower at higher latitudes, while normalised nestedness increased towards higher latitudes. Networks at lower latitudes in Europe are exposed to higher temperatures, which can result in higher visitation rates (Arroyo, Armesto, & Primack, 1985; Classen et al., 2015; Herrera, 2019), and the overall level of pollinator generalization is known to be higher at lower latitudes (Schleuning et al., 2012). These factors should increase the number of possible connections that can be established between plants and pollinators for a given network size, resulting in more connected and less nested networks at lower latitudes. Finally, one third of networks were more nested than expected by chance. While this supports the idea that plant-pollinator networks tend to be nested (Bascompte et al., 2003), this result also suggests that nestedness could be a less prevalent feature than previously thought for plant-pollinator networks (Payrató-Borras et al., 2019).

Despite this database covering a wide range of habitats across 23 countries, it contains geographical biases that can impact our understanding of plant-pollinator communities (Hughes et al., 2021). For instance, most plant-pollinator networks are sampled from central Europe, while Eastern Europe and the Mediterranean region are underrepresented. This is consistent with previous studies which also report lack of plant-pollinator data for those regions (Bennett et al., 2018; Marshall et al., 2024), highlighting that this database shows existing patterns in data availability despite the absence of a systematic search for studies. This lack of data is especially relevant for Eastern Europe which has vast landscapes of high quality semi-natural grasslands but is experiencing rapid land use change (Sutcliffe et al., 2015), and the Mediterranean region is likely to be severely impacted by climate change (Duchenne et al., 2020; Jaworski et al., 2022; Pareja-Bonilla, Arista, Morellato, & Ortiz, 2023). These areas are well known for their rich pollinator diversity (Miličić, Vujić, & Cardoso, 2018; Reverté et

al., 2023), and their under-representation is likely contributing to the low taxonomic coverage of this database at the European level. Although some of the most well studied countries in Europe (e.g., Belgium, The Netherlands) have already experienced land use change and biodiversity loss at the end of the 20th century (Carvalheiro et al., 2013), plant-pollinator communities in Europe and across the globe still face current and future threats from climate change (Bartomeus et al., 2011; Duchenne et al., 2020), land use change (Batáry, Dicks, Kleijn, & Sutherland, 2015; Reidsma, Tekelenburg, Van den Berg, & Alkemade, 2006), and the intro-duction of alien species (Vanbergen, Espíndola, & Aizen, 2018; Vilà et al., 2009). Therefore, continuous monitoring programs are needed in order to evaluate spatio-temporal changes of species and their interactions across different European habitats and regions. This will al-low local and large scale analyses of the status and trends of plant-pollinator communities, effectively informing management and conservation actions.

In conclusion, the EuPPollNet database enables researchers to explore spatial, taxonomic and structural properties of plant-pollinator networks within Europe. In contrast to previous databases, EuPPollNet provides interaction data along with sampling information that could help researchers to better control for sampling effort and completeness and to select the most suitable networks for their research questions. Here, we have shown how connectance and nest-edness change across their latitudinal range and that nestedness is not a ubiquitous feature of all plant-pollinator networks. These analyses aim to highlight the variability present across Europe in the structure of plant-pollinator networks and illustrate the opportunities available to develop and test questions about spatio-temporal network change using EuPPollNet. The reproducible workflow allows researchers to adapt and reuse this database, enabling the continuous addition of new networks to better evaluate the status and trends of plant-pollinator communities. Finally, we hope this database becomes an iterative resource that keeps growing and improving over time to better understand and conserve European biodiversity.

35 REFERENCES

- Almeida-Neto, M., Guimaraes, P., Guimaraes Jr, P. R., Loyola, R. D., & Ulrich, W. (2008).

 A consistent metric for nestedness analysis in ecological systems: Reconciling concept and measurement. *Oikos*, 117(8), 1227–1239.
- Archer, C. R., Pirk, C. W. W., Carvalheiro, L. G., & Nicolson, S. W. (2014). Economic and ecological implications of geographic bias in pollinator ecology in the light of pollinator declines. *Oikos*, 123(4), 401–407.
- Arroyo, M. T. K., Armesto, J. J., & Primack, R. B. (1985). Community studies in pollination ecology in the high temperate andes of central chile II. Effect of temperature on visitation rates and pollination possibilities. *Plant Systematics and Evolution*, 149, 187–203.
- Ballantyne, G., Baldock, K. C., & Willmer, P. G. (2015). Constructing more informative
 plant-pollinator networks: Visitation and pollen deposition networks in a heathland plant
 community. Proceedings of the Royal Society B: Biological Sciences, 282(1814), 20151130.
- Bartomeus, I., Ascher, J. S., Wagner, D., Danforth, B. N., Colla, S., Kornbluth, S., & Winfree,
 R. (2011). Climate-associated phenological advances in bee pollinators and bee-pollinated
 plants. Proceedings of the National Academy of Sciences, 108(51), 20645–20649.
- Bascompte, J., & Jordano, P. (2007). Plant-animal mutualistic networks: The architecture of biodiversity. *Annu. Rev. Ecol. Evol. Syst.*, 38, 567–593.
- Bascompte, J., Jordano, P., Melián, C. J., & Olesen, J. M. (2003). The nested assembly
 of plant–animal mutualistic networks. *Proceedings of the National Academy of Sciences*,
 100(16), 9383–9387.
- Batáry, P., Dicks, L. V., Kleijn, D., & Sutherland, W. J. (2015). The role of agri-environment
 schemes in conservation and environmental management. Conservation Biology, 29(4),
 1006–1016.
- Bellard, C., Leclerc, C., Leroy, B., Bakkenes, M., Veloz, S., Thuiller, W., & Courchamp,
 F. (2014). Vulnerability of biodiversity hotspots to global change. Global Ecology and
 Biogeography, 23(12), 1376–1386.
- Bennett, J. M., Thompson, A., Goia, I., Feldmann, R., Ştefan, V., Bogdan, A., et al.others. (2018). A review of european studies on pollination networks and pollen limitation, and a case study designed to fill in a gap. *AoB Plants*, 10(6), ply068.
- Burkle, L. A., & Alarcón, R. (2011). The future of plant–pollinator diversity: Understanding
 interaction networks across time, space, and global change. American Journal of Botany,
 98(3), 528–538.
- Carvalheiro, L. G., Kunin, W. E., Keil, P., Aguirre-Gutiérrez, J., Ellis, W. N., Fox, R., et
 al.others. (2013). Species richness declines and biotic homogenisation have slowed down
 for NW-european pollinators and plants. *Ecology Letters*, 16(7), 870–878.
- Chacoff, N. P., Vázquez, D. P., Lomáscolo, S. B., Stevani, E. L., Dorado, J., & Padrón, B.
 (2012). Evaluating sampling completeness in a desert plant–pollinator network. *Journal of Animal Ecology*, 81(1), 190–200.
- Chamberlain, S., Oldoni, D., & Waller, J. (2022). Rgbif: Interface to the global biodiversity
 information facility API.

- Classen, A., Peters, M. K., Kindeketa, W. J., Appelhans, T., Eardley, C. D., Gikungu, M. W.,

 577 ... Steffan-Dewenter, I. (2015). Temperature versus resource constraints: Which factors determine bee diversity on m ount k ilimanjaro, t anzania? Global Ecology and Biogeography,

 24(6), 642–652.
- Cribari-Neto, F., & Zeileis, A. (2010). Beta regression in r. *Journal of Statistical Software*, 34, 1–24.
- Culley, T. M., Weller, S. G., & Sakai, A. K. (2002). The evolution of wind pollination in angiosperms. *Trends in Ecology & Evolution*, 17(8), 361–369.
- Dormann, C. F., Gruber, B., & Fründ, J. (2008). Introducing the bipartite package: Analysing ecological networks. *Interaction*, 1(0.2413793), 8–11.
- Duchenne, F., Thébault, E., Michez, D., Elias, M., Drake, M., Persson, M., ... Fontaine, C.
 (2020). Phenological shifts alter the seasonal structure of pollinator assemblages in europe.
 Nature Ecology & Evolution, 4(1), 115–121.
- Eichenberg, D., Bowler, D. E., Bonn, A., Bruelheide, H., Grescho, V., Harter, D., ... Jansen, F.
 (2021). Widespread decline in central european plant diversity across six decades. Global
 Change Biology, 27(5), 1097–1110.
- Elberling, H., & Olesen, J. M. (1999). The structure of a high latitude plant-flower visitor system: The dominance of flies. *Ecography*, 22(3), 314–323.
- Encinas-Viso, F., Bovill, J., Albrecht, D. E., Florez-Fernandez, J., Lessard, B., Lumbers, J., ... Milla, L. (2023). Pollen DNA metabarcoding reveals cryptic diversity and high spatial turnover in alpine plant—pollinator networks. *Molecular Ecology*, 32(23), 6377–6393.
- García, Y., Giménez-Benavides, L., Iriondo, J. M., Lara-Romero, C., Méndez, M., Morente López, J., & Santamaría, S. (2024). Addition of nocturnal pollinators modifies the structure
 of pollination networks. Scientific Reports, 14(1), 1226.
- Gibson, R. H., Knott, B., Eberlein, T., & Memmott, J. (2011). Sampling method influences the structure of plant–pollinator networks. *Oikos*, 120(6), 822–831.
- Goulson, D., Nicholls, E., Botías, C., & Rotheray, E. L. (2015). Bee declines driven by combined stress from parasites, pesticides, and lack of flowers. *Science*, 347(6229), 1255957.
- Govaerts, R., Nic Lughadha, E., Black, N., Turner, R., & Paton, A. (2021). The world
 checklist of vascular plants, a continuously updated resource for exploring global plant
 diversity. Scientific Data, 8(1), 215.
- Grenié, M., Berti, E., Carvajal-Quintero, J., Dädlow, G. M. L., Sagouis, A., & Winter, M. (2023). Harmonizing taxon names in biodiversity data: A review of tools, databases and best practices. *Methods in Ecology and Evolution*, 14(1), 12–25.
- Guimaraes, P. R. (2020). The structure of ecological networks across levels of organization.
 Annual Review of Ecology, Evolution, and Systematics, 51, 433–460.
- Gusenbauer, M. (2019). Google scholar to overshadow them all? Comparing the sizes of 12 academic search engines and bibliographic databases. *Scientometrics*, 118(1), 177–214.
- Gusenbauer, M., & Haddaway, N. R. (2020). Which academic search systems are suitable for systematic reviews or meta-analyses? Evaluating retrieval qualities of google scholar, PubMed, and 26 other resources. Research Synthesis Methods, 11(2), 181–217.
- Hadfield, J. D. (2010). MCMC methods for multi-response generalized linear mixed models:

 The MCMCglmm r package. *Journal of Statistical Software*, 33, 1–22.

- Hass, A. L., Kormann, U. G., Tscharntke, T., Clough, Y., Baillod, A. B., Sirami, C., et
 al.others. (2018). Landscape configurational heterogeneity by small-scale agriculture, not
 crop diversity, maintains pollinators and plant reproduction in western europe. Proceedings
 of the Royal Society B: Biological Sciences, 285(1872), 20172242.
- Heberling, J. M., Miller, J. T., Noesgaard, D., Weingart, S. B., & Schigel, D. (2021). Data
 integration enables global biodiversity synthesis. Proceedings of the National Academy of
 Sciences, 118(6), e2018093118.
- Hedtke, S. M., Patiny, S., & Danforth, B. N. (2013). The bee tree of life: A supermatrix approach to apoid phylogeny and biogeography. *BMC Evolutionary Biology*, 13, 1–13.
- Herrera, C. M. (2019). Complex long-term dynamics of pollinator abundance in undisturbed mediterranean montane habitats over two decades. *Ecological Monographs*, 89(1), e01338.
- Herrera, C. M. (2020). Gradual replacement of wild bees by honeybees in flowers of the
 mediterranean basin over the last 50 years. Proceedings of the Royal Society B, 287(1921),
 20192657.
- 633 Hijmans, R. J., Bivand, R., Forner, K., Ooms, J., Pebesma, E., & Sumner, M. D. (2022).
 634 *Package "terra"*.
- Hoeppke, C., & Simmons, B. I. (2021). Maxnodf: An r package for fair and fast comparisons of nestedness between networks. *Methods in Ecology and Evolution*, 12(4), 580–585.
- Hsieh, T., & Chao, A. (2016). iNEXT: An r package for rarefaction and extrapolation of species diversity (h ill numbers). *Methods in Ecology and Evolution*, 7(12), 1451–1456.
- Hughes, A. C., Orr, M. C., Ma, K., Costello, M. J., Waller, J., Provoost, P., ... Qiao, H. (2021).
 Sampling biases shape our view of the natural world. *Ecography*, 44(9), 1259–1269.
- Hung, K.-L. J., Kingston, J. M., Albrecht, M., Holway, D. A., & Kohn, J. R. (2018). The
 worldwide importance of honey bees as pollinators in natural habitats. *Proceedings of the Royal Society B: Biological Sciences*, 285(1870), 20172140.
- Isaac, N. J., Cruickshanks, K. L., Weddle, A. M., Marcus Rowcliffe, J., Brereton, T. M., Dennis,
 R. L., ... Thomas, C. D. (2011). Distance sampling and the challenge of monitoring butterfly
 populations. Methods in Ecology and Evolution, 2(6), 585–594.
- Jaworski, C. C., Geslin, B., Zakardjian, M., Lecareux, C., Caillault, P., Nève, G., et al.others. (2022). Long-term experimental drought alters floral scent and pollinator visits in a mediter-ranean plant community despite overall limited impacts on plant phenotype and reproduction. *Journal of Ecology*, 110(11), 2628–2648.
- Jordano, P. (1987). Patterns of mutualistic interactions in pollination and seed dispersal:

 Connectance, dependence asymmetries, and coevolution. *The American Naturalist*, 129(5),
 657–677.
- Jordano, P. (2016). Sampling networks of ecological interactions. Functional Ecology, 30(12),
 1883–1893.
- Jordano, P., Bascompte, J., & Olesen, J. M. (2003). Invariant properties in coevolutionary networks of plant—animal interactions. *Ecology Letters*, 6(1), 69–81.
- Kleijn, D., Winfree, R., Bartomeus, I., Carvalheiro, L. G., Henry, M., Isaacs, R., et al.others.
 (2015). Delivery of crop pollination services is an insufficient argument for wild pollinator conservation. *Nature Communications*, 6(1), 7414.
- Leclercq, N., Marshall, L., Caruso, G., Schiel, K., Weekers, T., Carvalheiro, L. G., et al. others.

- (2023). European bee diversity: Taxonomic and phylogenetic patterns. *Journal of Biogeog*raphy, 50(7), 1244–1256.
- Lefebvre, V., Villemant, C., Fontaine, C., & Daugeron, C. (2018). Altitudinal, temporal and trophic partitioning of flower-visitors in alpine communities. *Scientific Reports*, 8(1), 4706.
- León-Osper, M., & Narbona, E. (2022). Unravelling the mystery of red flowers in the mediterranean basin: How to be conspicuous in a place dominated by hymenopteran pollinators. Functional Ecology, 36(11), 2774–2790.
- Li, D. (2023). Rtrees: An r package to assemble phylogenetic trees from megatrees. *Ecography*, 2023(7), e06643.
- Magrach, A., González-Varo, J. P., Boiffier, M., Vilà, M., & Bartomeus, I. (2017). Honeybee spillover reshuffles pollinator diets and affects plant reproductive success. *Nature Ecology Evolution*, 1(9), 1299–1307.
- Marshall, L., Leclercq, N., Carvalheiro, L. G., Dathe, H. H., Jacobi, B., Kuhlmann, M., ...
 Vereecken, N. J. (2024). Understanding and addressing shortfalls in european wild bee
 data. *Biological Conservation*, 290, 110455.
- Martínez-Núñez, C., Kleijn, D., Ganuza, C., Heupink, D., Raemakers, I., Vertommen, W., & Fijen, T. P. (2022). Temporal and spatial heterogeneity of semi-natural habitat, but not crop diversity, is correlated with landscape pollinator richness. *Journal of Applied Ecology*, 59(5), 1258–1267.
- Miličić, M., Vujić, A., & Cardoso, P. (2018). Effects of climate change on the distribution of
 hoverfly species (diptera: Syrphidae) in southeast europe. Biodiversity and Conservation,
 27, 1173–1187.
- Olesen, J. M., Bascompte, J., Dupont, Y. L., Elberling, H., Rasmussen, C., & Jordano, P. (2011). Missing and forbidden links in mutualistic networks. *Proceedings of the Royal Society B: Biological Sciences*, 278(1706), 725–732.
- Olesen, J. M., & Jordano, P. (2002). Geographic patterns in plant–pollinator mutualistic networks. *Ecology*, 83(9), 2416–2424.
- Ollerton, J. (2017). Pollinator diversity: Distribution, ecological function, and conservation.

 Annual Review of Ecology, Evolution, and Systematics, 48, 353–376.
- Orr, M. C., Hughes, A. C., Chesters, D., Pickering, J., Zhu, C.-D., & Ascher, J. S. (2021).
 Global patterns and drivers of bee distribution. *Current Biology*, 31(3), 451–458.
- Paradis, E., Blomberg, S., Bolker, B., Brown, J., Claude, J., Cuong, H. S., ... Didier, G. (2019).
 Package "ape." Analyses of Phylogenetics and Evolution, Version, 2(4), 47.
- Pareja-Bonilla, D., Arista, M., Morellato, L. P. C., & Ortiz, P. L. (2023). Better soon than never: Climate change induces strong phenological reassembly in the flowering of a mediterranean shrub community. *Annals of Botany*, mcad193.
- Payrató-Borras, C., Hernández, L., & Moreno, Y. (2019). Breaking the spell of nestedness:

 The entropic origin of nestedness in mutualistic systems. *Physical Review X*, 9(3), 031024.
- Pebesma, E. J. et al. (2018). Simple features for r: Standardized support for spatial vector data. R J., 10(1), 439.
- Poelen, J. H., Simons, J. D., & Mungall, C. J. (2014). Global biotic interactions: An open infrastructure to share and analyze species-interaction datasets. *Ecological Informatics*, 24, 148–159.

- Poisot, T., Baiser, B., Dunne, J. A., Kéfi, S., Massol, F., Mouquet, N., ... Gravel, D. (2016).
 Mangal-making ecological network analysis simple. *Ecography*, 39(4), 384–390.
- Poisot, T., Bergeron, G., Cazelles, K., Dallas, T., Gravel, D., MacDonald, A., ... Vissault, S.
 (2021). Global knowledge gaps in species interaction networks data. *Journal of Biogeogra-phy*, 48(7), 1552–1563.
- Proença, V., Martin, L. J., Pereira, H. M., Fernandez, M., McRae, L., Belnap, J., et al.others.
 (2017). Global biodiversity monitoring: From data sources to essential biodiversity variables. *Biological Conservation*, 213, 256–263.
- Ramos-Jiliberto, R., Domínguez, D., Espinoza, C., Lopez, G., Valdovinos, F. S., Bustamante, R. O., & Medel, R. (2010). Topological change of andean plant–pollinator networks along an altitudinal gradient. *Ecological Complexity*, 7(1), 86–90.
- Rech, A. R., Dalsgaard, B., Sandel, B., Sonne, J., Svenning, J.-C., Holmes, N., & Ollerton, J. (2016). The macroecology of animal versus wind pollination: Ecological factors are more important than historical climate stability. *Plant Ecology & Diversity*, 9(3), 253–262.
- Reidsma, P., Tekelenburg, T., Van den Berg, M., & Alkemade, R. (2006). Impacts of land-use change on biodiversity: An assessment of agricultural biodiversity in the european union.

 Agriculture, Ecosystems & Environment, 114(1), 86–102.
- Revell, L. J. (2012). Phytools: An r package for phylogenetic comparative biology (and other things). *Methods in Ecology and Evolution*, (2), 217–223.
- Reverté, S., Miličić, M., Ačanski, J., Andrić, A., Aracil, A., Aubert, M., et al.others. (2023).

 National records of 3000 european bee and hoverfly species: A contribution to pollinator conservation. *Insect Conservation and Diversity*, 16(6), 758–775.
- Sala, O. E., Stuart Chapin, F., Armesto, J. J., Berlow, E., Bloomfield, J., Dirzo, R., et al. others. (2000). Global biodiversity scenarios for the year 2100. Science, 287(5459), 1770–1774.
- Saunders, M. E., Kendall, L. K., Lanuza, J. B., Hall, M. A., Rader, R., & Stavert, J. R. (2023).
 Climate mediates roles of pollinator species in plant–pollinator networks. Global Ecology
 and Biogeography, 32(4), 511–518.
- Schleuning, M., Fründ, J., Klein, A.-M., Abrahamczyk, S., Alarcón, R., Albrecht, M., et
 al.others. (2012). Specialization of mutualistic interaction networks decreases toward tropical latitudes. Current Biology, 22(20), 1925–1931.
- Schwarz, B., Vázquez, D. P., CaraDonna, P. J., Knight, T. M., Benadi, G., Dormann, C. F., et al. others. (2020). Temporal scale-dependence of plant–pollinator networks. *Oikos*, 129(9), 1289–1302.
- Settele, J., Bishop, J., & Potts, S. G. (2016). Climate change impacts on pollination. Nature Plants, 2(7), 1–3.
- Simpson, D. T., Weinman, L. R., Genung, M. A., Roswell, M., MacLeod, M., & Winfree, R. (2022). Many bee species, including rare species, are important for function of entire plant-pollinator networks. *Proceedings of the Royal Society B*, 289(1972), 20212689.
- Smith, S. A., & Brown, J. W. (2018). Constructing a broadly inclusive seed plant phylogeny.

 American Journal of Botany, 105(3), 302–314.
- Song, C., Rohr, R. P., & Saavedra, S. (2017). Why are some plant–pollinator networks more nested than others? *Journal of Animal Ecology*, 86(6), 1417–1424.
- Staniczenko, P., Kopp, J. C., & Allesina, S. (2013). The ghost of nestedness in ecological

- networks. Nature Communications, 4(1), 1–6.
- 749 Stanners, D., & Bourdeau, P. (1995). Europe's environment: The dobříš assessment.
- Steffan-Dewenter, I., & Tscharntke, T. (2000). Resource overlap and possible competition between honey bees and wild bees in central europe. *Oecologia*, 122, 288–296.
- Sutcliffe, L. M., Batáry, P., Kormann, U., Báldi, A., Dicks, L. V., Herzon, I., et al.others.
 (2015). Harnessing the biodiversity value of central and eastern european farmland. *Diversity and Distributions*, 21(6), 722–730.
- Thompson, G. G., & Withers, P. C. (2003). Effect of species richness and relative abundance on the shape of the species accumulation curve. *Austral Ecology*, 28(4), 355–360.
- Traveset, A., Tur, C., Trøjelsgaard, K., Heleno, R., Castro-Urgal, R., & Olesen, J. M. (2016).
 Global patterns of mainland and insular pollination networks. *Global Ecology and Biogeography*, 25(7), 880–890.
- Troia, M. J., & McManamay, R. A. (2016). Filling in the GAPS: Evaluating completeness and
 coverage of open-access biodiversity databases in the united states. *Ecology and Evolution*,
 6(14), 4654–4669.
- Trøjelsgaard, K., & Olesen, J. M. (2013). Macroecology of pollination networks. Global Ecology and Biogeography, 22(2), 149–162.
- Van der Loo, M. P. et al. (2014). The stringdist package for approximate string matching. R J., 6(1), 111.
- Vanbergen, A. J., Espíndola, A., & Aizen, M. A. (2018). Risks to pollinators and pollination from invasive alien species. *Nature Ecology & Evolution*, 2(1), 16–25.
- Vilà, M., Bartomeus, I., Dietzsch, A. C., Petanidou, T., Steffan-Dewenter, I., Stout, J. C., &
 Tscheulin, T. (2009). Invasive plant integration into native plant–pollinator networks across
 europe. Proceedings of the Royal Society B: Biological Sciences, 276(1674), 3887–3893.
- Vizentin-Bugoni, J., Maruyama, P. K., Souza, C. S. de, Ollerton, J., Rech, A. R., & Sazima,
 M. (2018). Plant-pollinator networks in the tropics: A review. Ecological Networks in the
 Tropics: An Integrative Overview of Species Interactions from Some of the Most Species Rich Habitats on Earth, 73–91.
- Wetzel, F. T., Bingham, H. C., Groom, Q., Haase, P., Kõljalg, U., Kuhlmann, M., et al.others.
 (2018). Unlocking biodiversity data: Prioritization and filling the gaps in biodiversity observation data in europe. *Biological Conservation*, 221, 78–85.
- Wiemers, M., Balletto, E., Dincă, V., Fric, Z. F., Lamas, G., Lukhtanov, V., et al.others.
 (2018). An updated checklist of the european butterflies (lepidoptera, papilionoidea).
 ZooKeys, (811), 9.
- Windsor, F. M., Hoogen, J. van den, Crowther, T. W., & Evans, D. M. (2023). Using ecological networks to answer questions in global biogeography and ecology. *Journal of Biogeography*, 50(1), 57-69.
- Winfree, R., Reilly, J. R., Bartomeus, I., Cariveau, D. P., Williams, N. M., & Gibbs, J. (2018).
 Species turnover promotes the importance of bee diversity for crop pollination at regional scales. Science, 359(6377), 791–793.
- Zattara, E. E., & Aizen, M. A. (2021). Worldwide occurrence records suggest a global decline in bee species richness. *One Earth*, 4(1), 114–123.

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801 CONFLICT OF INTEREST

802 None.

803 DATA AVAILABILITY

All data and code to produce of this database and manuscript are available at Zenodo (LINK) and Github (https://github.com/JoseBSL/EuPPollNet.git).

506 SUPPORTING INFORMATION

Title: EuPPollNet: A European database of plant-pollinator networks

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833 Contains:

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- Supplementary text 1
- Figure S1
- Figure S2
- Figure S3

Supplementary text 1

Habitat definitions: 839

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- 1) Ruderal vegetation: Plants growing on highly disturbed sites such as road sides or 840 mineral extraction sites. 841
- 2) Agricultural margins: Sides of crops that can include any type of vegetation from low 842 growing plants to trees. 843
- 3) Green urban areas: Parks, private gardens or small pastures within an urban setting. Botanical gardens are included in this category. 845
- 4) Agricultural land: Includes any type of crop and any type of vegetation growing within 846 them. 847
 - 5) Forest/woodland understory: Any plant community sampled under a wooded group of plants. The forest could be embedded in an agricultural setting or in a fully natural scenario. We have included in this category agro-forestry areas and open to dense forest. Note that we have excluded from this category forest that contains sclerophyllous vegetation.
 - 6) Semi-natural grassland: Low growing plant community with relatively low disturbances but under low pressure such as seasonal moving or extensive grazing.
- 7) Intensive grassland: Any type of low growing plant community that is highly in-855 fluenced by human disturbance. For instance, agriculture, mowing, moderate to high 856 grazing or urban environments. Note that this category also includes old pastures with 857 regrowth of woody vegetation. 858
 - 8) Sclerophyllous vegetation: Any type of system with a dominant shrub community adapted to drought. Typical of the Mediterranean region. Note, that we have included in this category also woodlands (open coniferous forest) where the shrub community was the main focus of the study.
 - 9) Beaches, dunes, sands: Plant communities growing on sandy soil.
 - 10) Riparian vegetation: Plant communities growing on river margins.
- 11) Alpine grasslands: Low growing plant communities with little or none human distur-865 bance. Often located at high elevation areas within Europe. 866
- 12) Moors and heathland: Low growing woody vegetation characteristic from low fertile 867 soils near the coast or in alpine areas. 868

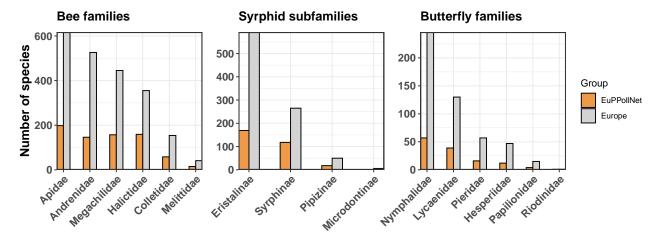


Figure S1. Coverage of the EuPPollNet species for bees (family level), syrphids (subfamily level) and butterflies (family level) in relation to the total number of European species within these taxonomic groups.

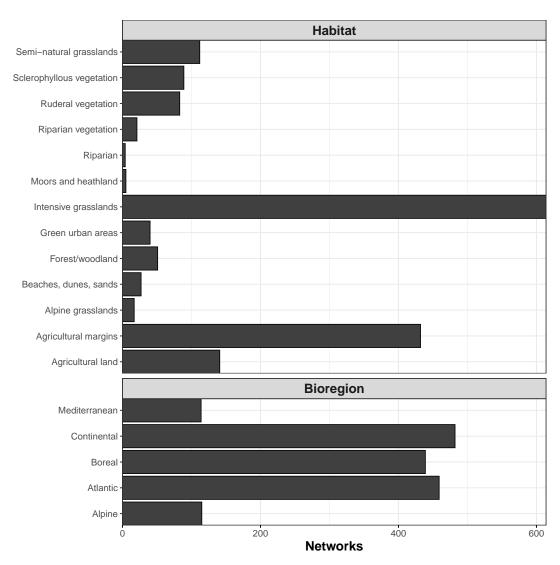


Figure S2. Barplot indicating the number of networks by habitat and bioclimatic region within the database.

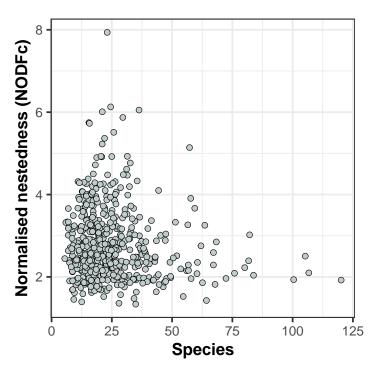


Figure S3. Association between nestedness (NODFc) and the geometric mean of plant and pollinator species per network