

DATA ARTICLE

The atlas of mitochondrial genetic diversity for Western Palaearctic butterflies

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Abstract

Motivation: Butterflies represent a model in biology and a flagship group for invertebrate conservation. We provide four new resources for the Western Palaearctic butterflies: (1) an updated checklist comprising 552 species; (2) a curated dataset of 32,126 mitochondrial cytochrome c oxidase subunit I (COI) sequences for 532 species, including a *de novo* reference library for the Maghreb (Morocco, northern Algeria and Tunisia) and Macaronesia (Azores, Madeira and Canary Islands); (3) seven indexes of intraspecific genetic variation (IGV): observed and expected number of haplotypes, haplotype and nucleotide diversity, two fixation indexes and maximum p-distance; and (4) species-level maps illustrating the distribution of COI variability and haplotype networks. The updated checklist will be fundamental for any application dealing with butterfly diversity in the Western Palaearctic. The IGV indexes provide measures for genetic polymorphism and spatial structure and represent proxies for dispersal capacity. These resources will facilitate comparative studies of macrogenetics, foster integrative taxonomy and aid conservation strategies.

Main types of variables contained: A complete species checklist in table format, 32,126 mitochondrial DNA barcodes provided with metadata (species membership, WGS84 coordinates and sequence length) and a book in PDF format, including the IGV atlas and indexes, are provided.

Spatial location and grain: The checklist encompasses Europe up to the Urals in the east, north Macaronesia (the Azores, Madeira and the Canary Islands) and the Maghreb (Morocco, northern Algeria and Tunisia). COI sequences have been retained in the geographical interval of -31.3 to 67.5° of longitude and 27.5 – 71.2° of latitude.

Time period and grain: COI sequences originate from studies published between 1998 and 2022 and from *de novo* sequencing of 2541 specimens done between 2007 and 2022.

Leonardo Dapporto and Mattia Menchetti contributed equally as first authors.

Vlad Dincă and Roger Vila contributed equally as senior authors.

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Major taxa and level of measurement: Butterflies (Lepidoptera: Papilionoidea), analysed from individual to species level.

Software format: Data and functions to manage the dataset are provided in the ioda-tabase R package (<https://github.com/leondap/iodatabase>) and in Dryad (<https://doi.org/10.5061/dryad.9w0vt4bjj>).

KEYWORDS

COI sequences, DNA barcoding library, intraspecific genetic variation, Papilionoidea, phylogeography

1 | INTRODUCTION

The advent of phylogeography (Avise et al., 1987) sparked the commitment of mapping the fundamental layer of biodiversity encompassed by intraspecific genetic variation (IGV). Nevertheless, mapping IGV remained a vision for decades. Recently, faster and more affordable DNA sequencing has enabled a rapid accumulation of data for a larger proportion of biodiversity. Repositories, such as GenBank and BOLD, curate $>10^6$ cytochrome c oxidase subunit I (COI) sequences (the standard mitochondrial marker for animal DNA barcoding) (Hebert et al., 2003), in many cases complemented by high-quality metadata. As COI sequences accumulate, comparative macrogenetic analyses become possible (Leigh et al., 2021; Theodoridis et al., 2020).

Here, we have gathered and improved large-scale biodiversity surveys to generate an IGV atlas for Western Palaearctic butterflies. The primary resources we referred to are as follows: (1) the checklists of Western Palaearctic butterflies by Wiemers et al. (2018) and Middleton-Welling et al. (2020); and (2) a series of DNA-barcode libraries of Eurasiatic butterflies (Dapporto et al., 2019; Dincă et al., 2011, 2015, 2021; Hausmann et al., 2011; Huemer & Tarmann, 2016; Litman et al., 2018; Lukhtanov et al., 2009; Menchetti et al., 2021). We revised the checklists and added 2541 specimens sequenced *de novo*, including a comprehensive DNA-barcode library for Maghreb and Macaronesia. We provided COI sequences and IGV assessments for 532 species at the crossroads of three continents (Africa, Europe and Asia) and compiled a complete set of networks and distribution maps for haplotypes. To our knowledge, this is the first attempt to build a complete IGV atlas at continental and inter-continental levels. We hope that this resource will pilot similar studies on other clades and/or regions and will represent a milestone in the identification, description and protection of hidden diversity.

2 | METHODS

2.1 | Checklists and geographical ranges

We revised and updated a recent butterfly checklist for Europe (Wiemers et al., 2018) and one for Europe and Maghreb (Middleton-Welling et al., 2020) (Table S1). For the taxonomic revision, we

accepted the conclusions of the most updated literature combining at least quantitative morphology and different genetic markers or based on genomic approaches (including double digest restriction-site associated DNA sequencing and phylogenomics) (see Table S2 and references therein). When evidence was insufficient, we refrained from applying changes, except in cases of deeply paraphyletic COI trees, which we treated as separate species.

We defined the Western Palaearctic “taxonomic area” (blue line in Figure 1) as follows. In the west, we included the Azores, Madeira and the Canary Islands. To delimit the fringes of the Sahara Desert, we set the southern limit in Morocco at 27.5°, followed the political border of this country to the north-east and set the southern boundary in Algeria and Tunisia at 33°. We included the European mainland and the islands under European administration in the Mediterranean, North and Baltic seas. No obvious faunistic boundaries exist to the east, and following previous checklists (Middleton-Welling et al., 2020; Wiemers et al., 2018) we included the western slopes of the Urals and excluded the Caucasus and western Kazakhstan. Given that any geographical boundary includes taxa at their distribution limit, we used a wider “dataset area” to provide context (Figure 1). Thus, we collected COI sequences of species in our checklist from the area included within the coordinates of −31.3° longitude (Flores, Azores), 27.5° latitude (El Hierro, the Canary Islands), 67.5° longitude (easternmost Urals) and 71.2° latitude (North Cape).

2.2 | Data acquisition, curation and quality control

We gathered COI sequences (DNA-barcode region between 400 and 658 bp) from three main sources: (1) *de novo* sequencing using standard procedures (deWaard et al., 2008) for most of the Maghreb and Macaronesia, but also improving coverage in Europe; (2) published DNA-barcode libraries compiled at the regional level (Dapporto et al., 2019; Dincă et al., 2011, 2015, 2021; Hausmann et al., 2011; Huemer & Tarmann, 2016; Litman et al., 2018; Menchetti et al., 2021) and other publicly available BOLD projects (e.g., LON-NorBOL for Norway); and (3) studies providing COI of single species or genera (references are available in Appendix S1). In the last case, we checked whether haplotypes, instead of specimens, were included in repositories (Paz-Vinas et al., 2021). If the number of specimens sharing a given haplotype in a particular location was reported

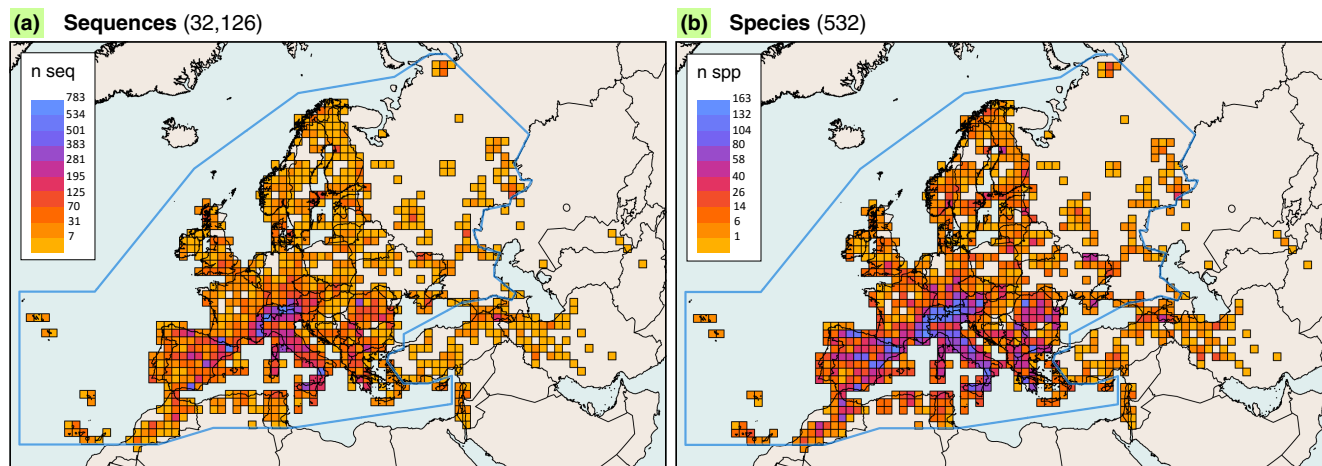


FIGURE 1 The study area with the representation of the “taxonomic area” (blue perimeter; Middleton-Welling et al., 2020) from where a complete species checklist has been assessed and updated. COI sequences of species also occurring outside the taxonomic area are included. (a) Number of sequences and (b) number of species for each 100 km × 100 km square cell. Variation in the number of DNA barcodes largely reflects butterfly richness across the taxonomic area.

(Ehl et al., 2021; Zinetti et al., 2013), we replicated the haplotype sequences to obtain data at the specimen level; otherwise, the sequences were excluded.

We verified species identifications by building neighbour-joining trees for each genus. When morphology could not be verified, we removed sequences not clustering within the species to which they were attributed if the mismatch did not involve one of the 74 species showing DNA-barcode sharing (updated from Dincă et al., 2021) (Table S3).

Coordinates (in decimal degrees) were retrieved from meta-data and converted to the World Geodetic System 1984 (WGS84). When only an identifiable toponym was provided, we obtained approximate WGS84 coordinates at a resolution of 0.1° latitude and longitude. To obtain mainland areas of similar size, WGS84 coordinates were transformed into the Lambert azimuthal equal-area projection.

2.3 | Indexes of genetic variation

Based on COI sequences, we computed fundamental indexes of IGV. Initially, we obtained the number of haplotypes per species with the *haplotype* function of the *pegas* R package (Paradis, 2010). Then, for species with ≥10 sequences, we obtained the expected (asymptotic) number of haplotypes in the study area by rarefaction curves (D'Ercole et al., 2021; Dincă et al., 2021) using the *iNEXT* R package (Chao et al., 2014). Using the *hap.div* and *nuc.div* functions (*pegas*), we also calculated haplotype diversity (the probability that two sequences belong to the same haplotype) and nucleotide diversity (the average number of nucleotide differences per site between sequences). We calculated pairwise p-distances (the number of substitutions per site with pairwise deletion of missing sites) between sequences for each species and recorded the maximum p-distance among sequences.

The DST and GST fixation indexes (Meirmans & Hedrick, 2011; Nei, 1973) are defined as $(HT - HS)$ and DST/HT , respectively, where HT is the mean p-distance among all sequences and HS is the mean p-distance among sequences within the same populations (defined below). DST varies between negative values and the maximum p-distance. Negative values (intrapopulation differentiation higher than average differentiation) often represent artefacts attributable to relatively small sample sizes and are usually set to zero (Meirmans & Hedrick, 2011), thus we applied this solution.

GST is a standardized dimensionless index ranging between negative values (set to zero) and one, with zero implying panmixia and one a complete spatial structuring. In species showing a single haplotype, DST is zero and GST is not defined. We also set GST to zero in these cases because, in the absence of any IGV, population structure can be argued to be absent.

Computing fixation indexes requires a division of specimens by populations, and the division affects the results. To minimize this effect, records were binned to different sets of areas identified by islands and by mainland grid cells of 100 km × 100 km, 200 km × 200 km, 300 km × 300 km, 400 km × 400 km and 500 km × 500 km. Furthermore, we replicated the five analyses by placing the centre of each cell on the vertex among cells in the previous division. We obtained 10 different partitions and, after computing DST and GST for each of them, we averaged their values to obtain mean indexes. Fixation indexes were computed for species having ≥10 specimens included in at least two areas with no less than three specimens each. This minimum requirement appeared to be sufficiently reliable in DST and GST computation (Scalcio et al., 2020).

Nucleotide diversity and GST are of particular interest because they express the average divergence among sequences and how much different haplotypes are segregated in different areas, respectively. For this reason, we plotted these values on bivariate bubble plots showing the distribution of GST and nucleotide diversity

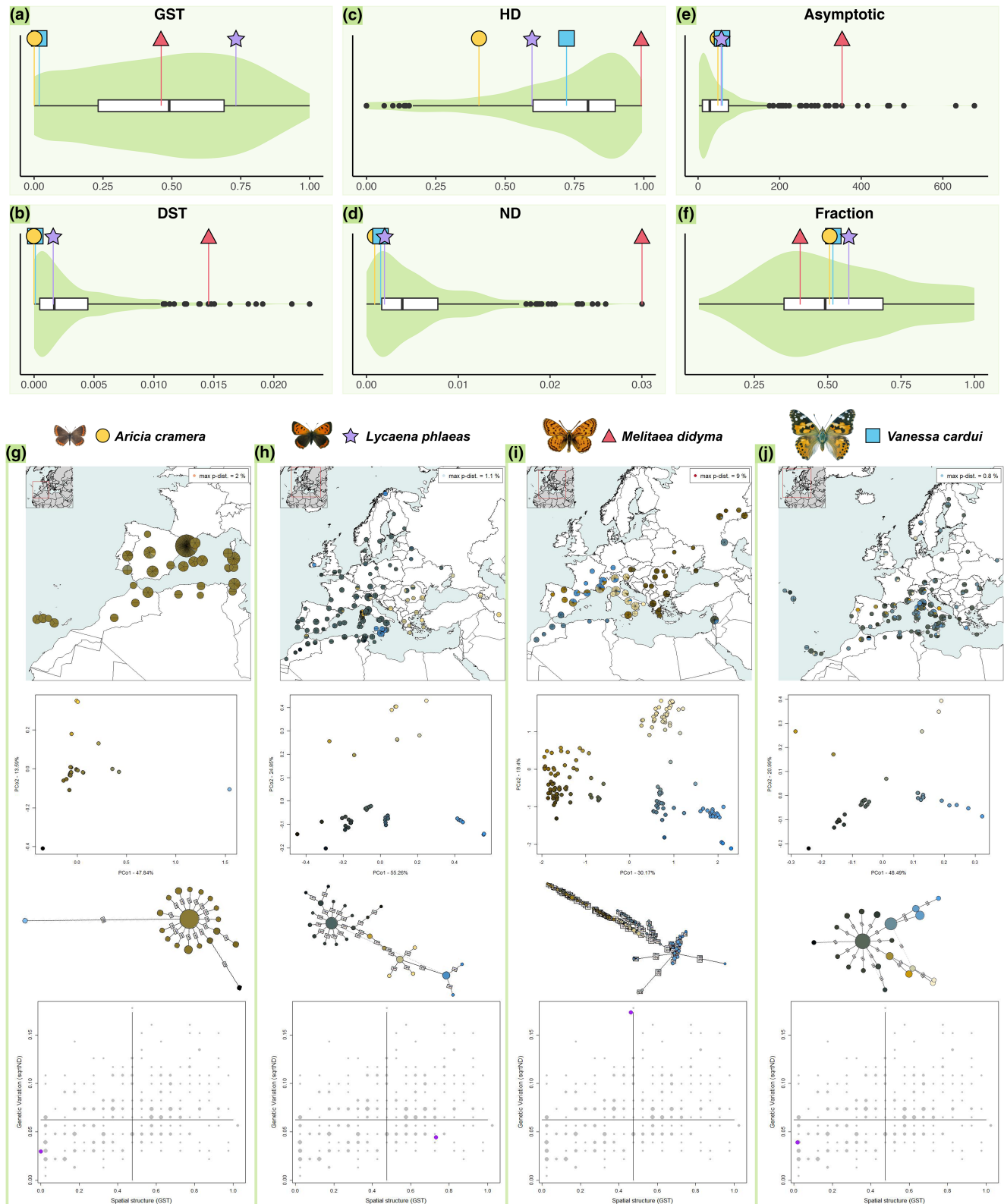


FIGURE 2 The distribution of intraspecific genetic variation (IGV) indexes: (a) GST; (b) DST; (c) haplotype diversity (HD); (d) nucleotide diversity (ND); (e) asymptotic (expected) number of haplotypes per species; (f) fraction of observed on expected number of haplotypes; and (g–j) four paradigmatic examples of the atlas representations with (from top to bottom): pie charts of sequenced specimens; the principal coordinates analysis (PCoA) in red–green–blue (RGB) space; haplotype networks, with colours for haplotypes as in pie charts, and scatterplots representing the GST and the square root-transformed values of nucleotide diversity with respect to all species in the dataset (grey bubbles with size proportional to the number of species in the intervals). The symbols for species in g–j are used to indicate their values in a–f.

(square root transformed) and their median values for all species in the dataset.

2.4 | Maps of genetic variation

We generated maps of haplotype distribution for each species with pie charts by attributing colours to sequences according to their p-distances. For each species, we subjected the p-distances matrix to a principal coordinates analysis (PCoA). We placed the two-dimensional configuration over a square having the colours yellow, blue, white and black at the corners and all possible shades in between, such that genetic similarity between two haplotypes was mirrored by colour similarity. The palette has been tested for perception by people with the most common types of colour blindness (Figure S4 in Appendix S1). The sequences were illustrated in pies grouped by areas of the same size; we used a variable cell size depending on the plotted area obtained by summing 0.3 to 1‰ of the latitudinal extension up to a maximum of 300 km × 300 km.

2.5 | Haplotype networks

For the construction of haplotype networks, we used sequences ≥600 bp. Haplotype networks were calculated with the randomized minimum spanning tree algorithm (Paradis, 2018) implemented in the *rmst* function of the R package *pegas*. Numbers of mutations among haplotypes were indicated using numbers over connections, and the haplotypes were assigned the colours obtained after PCoA.

2.6 | Script availability

The updated species checklist, COI sequences, metadata, the atlas and R scripts are available in Dryad (<https://doi.org/10.5061/dryad.9w0vt4bjj>), in the iodb database R package designed to store and manage the dataset (<https://github.com/leondap/iodb>) and in the DS-ATLAS (<https://doi.org/10.5883/DS-ATLAS>), a BOLD dataset where the newly sequenced specimens are released publicly. The atlas is also available in Appendix S1.

3 | RESULTS AND DISCUSSION

We updated the latest butterfly checklists, adding 16 species and excluding four taxa recently recognized as subspecies (Table S1). Based on this checklist (Table S2), comprising 552 species present in the “taxonomic area”, we obtained 32,126 geo-referenced COI sequences ≥400 bp from specimens within the “dataset area”, including 2541 previously unpublished sequences. A total of 532 species (96.4% of the checklist) were represented, with a mean of 58.2 (±88.0 SD) sequences per species.

The mean number of haplotypes per species was 16.2 (±20.8 SD) (for IGV indexes, see Table S2). As observed by Dincă et al. (2021), the number of haplotypes detected was often much lower than the asymptotic value (mean ± SD completeness 52.8 ± 23.6%), owing to the occurrence of few common haplotypes and many rare ones, frequently differentiated by only one or few mutations. Accordingly, haplotype diversity was skewed towards high values (many haplotypes), whereas nucleotide diversity was skewed towards low values (low divergences) (Figure 2).

GST showed an almost normal distribution, whereas DST was skewed towards low differentiation. The species showing low GST are characterized by very low haplotype and nucleotide diversity or by relatively high haplotype and nucleotide diversity with almost complete panmixia (Figure 2a–d). Conversely, some species with low haplotype diversity showed high GST when haplotypes exhibited strong spatial structure (Figure 2h).

The maps, which are provided in a supplementary book available as a main resource of this study (Appendix S1), revealed a variety of patterns. A strong divergence between continental areas (the Maghreb, Europe and the Middle East) emerged as a major pattern (e.g., *Aporia crataegi*, *Melitaea cinxia* and *Chazara briseis*). Many species (e.g., *Thymelicus lineola*, *Pyrgus serratalae* and *Lasiommata maera*) also showed differentiation among glacial refugia in southern Europe (Iberian, Italian and Balkan peninsulas), in line with the paradigm of the genetic legacy of ice ages on European biota (Hewitt, 2000; Taberlet et al., 1998).

This dataset facilitates in-depth comparative assessments by exploring general trends in distribution patterns, although this lies beyond the aim of this paper. Such an approach is essential to avoid potential cases of pseudocongruence that occur when researchers attempt to categorize patterns visually (Soltis et al., 2006). It also represents a tool for biologists studying the butterfly species covered, because it can be a great help in the planning of sampling intended to describe IGV further, allows for testing potential eco-evolutionary correlates, and highlights potential cryptic species or other evolutionary phenomena (e.g., introgression and genetic sweeps) to be studied in greater detail. Although mitochondrial IGV can have limited power in the inference of demographic processes and encompass only female patterns (Schmidt & Garraway, 2021), indexes of mitochondrial DNA variation are used as proxies for dispersal capacity (e.g., Melero et al., 2022). When enough data for nuclear genomes become available, it will be possible to clarify whether spatial patterns reflected in both mitochondrial and nuclear genomes are generally comparable. Finally, the allopatric/parapatric haplogroups documented here could help in defining subspecific taxonomy and can be viewed as management units for conservation.

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

DATA AVAILABILITY STATEMENT

The R scripts and COI sequences are available in Dryad (<https://doi.org/10.5061/dryad.9wOvt4bjj>) and in the iodatabase GitHub page (<https://github.com/leonardap/iodatabase>), where the FASTA files and the metadata comprising BOLD process IDs and GenBank accession numbers can be found. An R package (iodatabase) can also be installed from Github (<https://github.com/leonardap/iodatabase>), where a detailed description of the new functions is provided. The atlas of mitochondrial genetic diversity for Western Palaearctic butterflies is available as a PDF file in the same repositories and as Appendix S1.

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BIOSKETCH

Members of the research team are actively engaged in insect biogeography, systematics and conservation, with particular interest in analysing large-scale data generated by occurrence records and DNA sequencing. The main aim of their research is unravelling the historical and present-day factors responsible for species distributions across the Western Palearctic region.

SUPPORTING INFORMATION

Additional supporting information can be found online in the Supporting Information section at the end of this article.

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