Entomological Science



Functional groups of hoverflies in Southeast Europe across different vegetation types

Journal:	Entomological Science				
Manuscript ID	ENS-2020-0183.R2				
Wiley - Manuscript type:	Original Article				
Date Submitted by the Author:	n/a				
Complete List of Authors:	Miličić, Marija; University of Novi Sad, BioSense Institute – Research Institute for Information Technologies in Biosystems, Centre for Biosystems Popov, Snežana; University of Novi Sad Faculty of Science and Mathematics, Department of Biology and Ecology Jurca, Tamara; University of Novi Sad Faculty of Science and Mathematics, Department of Biology and Ecology Cardoso, Pedro; Finnish Museum of Natural History Janković, Marina; University of Novi Sad Faculty of Science and Mathematics, Department of Biology and Ecology Cardoso, Pedro; Finnish Museum of Natural History Janković, Marina; University of Novi Sad Faculty of Science and Mathematics, Department of Biology and Ecology Ačanski, Jelena; University of Novi Sad, BioSense Institute – Research Institute for Information Technologies in Biosystems, Centre for Biosystems Vujić, Ante; University of Novi Sad Faculty of Science and Mathematics, Department of Biology and Ecology				
Keywords:	Diptera, functional classification, insects, plant cover, richness, Syrphidae, traits				
Abstract:	To better understand the relationship between biodiversity and ecosystem functioning, it is increasingly accepted that the focus of study needs to shift from taxonomic identity to the diversity of functional traits displayed by species within a community. Such an approach allows species to be grouped according to particular functional characteristics. Increasingly viewed as an extremely important group of model organisms, hoverflies have been the focus of a variety of ecological studies. Based on data regarding selected functional traits of hoverflies registered in Southeast Europe, the main aims of our study were to define hoverfly functional groups according to the similarity of these traits, as well as to compare the representation of delineated hoverfly functional groups among these vegetation types. We used fuzzy clustering to classify 568 SE European hoverfly species into five functional groups. The principle trait separating these functional groups was larval feeding type, followed by size of species range, flight ability, number of generations, inundation tolerance, and tolerance to human impact. For 9 of 11 vegetation types, the dominant functional group was characterized by species with good flight ability, having high human impact tolerance and more annual generations. The remaining two vegetation types, South-west Balkan sub-Mediterranean mixed oak				

1	
2	
3 4 5 6 7 8	forests and Mediterranean mixed forests, showed disparate dominance patterns, indicating that richness of functional groups is dependent on vegetation. Further investigation of whether and how established conservation measures enable recovery of the functional richness affected by habitat disturbance would help elucidate the importance of functional diversity in preserving biodiversity.
9	
10 11	
12	
13	SCHOLAR ONE [™]
14	Manuscripts
15	
10	
18	
19	
20	
21	
23	
24	
25	
26 27	
27	
29	
30	
31	
32 33	
34	
35	
36	
3/	
39	
40	
41	
42	
45 44	
45	
46	
47	
48 49	
50	
51	
52	
53	
54 55	
56	
57	
58	
59 60	
00	

Functional groups of hoverflies in Southeast Europe across different vegetation types

Marija MILIČIĆ^{1,2*}, Snežana POPOV³, Tamara JURCA³, Pedro CARDOSO², Marina
 JANKOVIĆ³, Jelena AČANSKI¹, Ante VUJIĆ³

¹University of Novi Sad, BioSense Institute – Research Institute for Information Technologies in Biosystems, Dr
Zorana Đinđića 1, 21000 Novi Sad, Serbia: marija.milicic@biosense.rs; acanski@biosense.rs, ²LIBRe – Laboratory
for Integrative Biodiversity Research, Finnish Museum of Natural History, University of Helsinki, PO17 (Pohjoinen
Rautatiekatu 13), 00014, Helsinki, Finland: pedro.cardoso@helsinki.fi, ³University of Novi Sad, Faculty of
Sciences, ³Department of Biology and Ecology, Trg Dositeja Obradovića 2, Novi Sad, Serbia:
snezana.jovicic@dbe.uns.ac.rs; tamara.jurca@dbe.uns.ac.rs; marinaj@dbe.uns.ac.rs; ante.vujic@dbe.uns.ac.rs

*Correspondence: Marija Miličić, BioSense Institute – Research Institute for Information Technologies in
 Biosystems, University of Novi Sad, Dr Zorana Đinđića 1, 21000 Novi Sad, Serbia

13 Email: marija.milicic@biosense.rs

15 Abstract

To better understand the relationship between biodiversity and ecosystem functioning, it is increasingly accepted that the focus of study needs to shift from taxonomic identity to the diversity of functional traits displayed by species within a community. Such an approach allows species to be grouped according to particular functional characteristics. Increasingly viewed as an extremely important group of model organisms, hoverflies have been the focus of a variety of ecological studies. Based on data regarding selected functional traits of hoverflies registered in Southeast Europe, the main aims of our study were to define hoverfly functional groups according to the similarity of these traits, as well as to compare the representation of delineated hoverfly functional groups among these vegetation types. We used fuzzy clustering to classify 568 SE European hoverfly species into five functional groups. The principle trait separating these functional groups was larval feeding type, followed by size of species range, flight ability, number of generations, inundation tolerance, and tolerance to human impact. For 9 of 11 vegetation types, the dominant functional group was characterized by species with good flight ability, having high human impact tolerance and more annual generations. The remaining two vegetation types, South-west Balkan sub-Mediterranean mixed oak forests and Mediterranean mixed forests, showed disparate dominance patterns, indicating that richness of functional groups is dependent on vegetation. Further investigation of whether and how established conservation measures enable recovery of the functional richness affected by habitat disturbance would help elucidate the importance of functional diversity in preserving biodiversity.

35 Key words: Diptera, functional classification, insects, plant cover, richness, Syrphidae, traits

INTRODUCTION

Page 3 of 26

Entomological Science

Species richness and abundance have commonly been used as indicators to evaluate the state of a given ecosystem or ecosystem process (Medellín *et al.* 2000; Peters *et al.* 2016). However, a diverse and species-rich community does not necessarily mean that ecosystem functions or services are intact and function properly (Winsa *et al.* 2017). To better understand the relationship between biodiversity and (ecosystem) functioning, it is increasingly accepted that our focus needs to shift from taxonomic identity to the diversity of functional traits exhibited by species within a community (Díaz & Cabido 2001; Cadotte *et al.* 2011; Cardinale *et al.* 2012).

A functional group can be defined as a set of species displaying a similar response to the environment or having similar effects on ecosystem processes (Gitay & Noble 1997). Functional classification often has two objectives, one being to investigate the effects of species on ecosystem characteristics (Cornwell & Ackerly 2009), and the other to explore the type of response to environmental changes (functional response groups), such as habitat loss or degradation (Craven et al. 2016), availability of resources (Perkins et al. 2018), or climate change (Ooi et al. 2014). Identification of functional response groups may help to understand and predict how certain aspects of the community and ecosystem can be affected by environmental changes (Hooper et al. 2002).

Hoverflies are increasingly viewed as an extremely important group of model organisms with remarkable ecological and cultural value as pollinators (Jauker & Wolters 2008; Doyle et al. 2020) and biological control agents (Grosskopf 2005; Day et al. 2015), and they have been the focus of diverse ecological studies. Considerable effort has been devoted to investigating how hoverflies respond to the biggest environmental challenges worldwide, such as intensive agriculture (Li et al. 2020), urbanization (Persson et al. 2020), climate change (Miličić et al. 2018), and invasive species (Davis et al. 2018; Szigeti et al. 2020). However, the majority of past research has explored links between hoverfly species richness, abundance and/or distribution patterns, and environmental conditions, but very few studies have considered evaluating hoverfly functional groups (Schweiger et al. 2007; Keil et al. 2008).

Southeast Europe (SE Europe: Balkan Peninsula and the Aegean islands) is a region rich in flora (Sabovljević et al. 2008) and fauna (Crnobrnja-Isailović 2007; Poulakakis et al. 2015). Its geographical position at a crossroad of biogeographic influences, reliefs, climatic types and underlying bedrock preconditions it for high biodiversity within a relatively small area (Sabovljević et al. 2008). This region has been designated as one of the world's hotspots for hoverflies, harboring a great number of endemic and rare species (Vujić *et al.* 2001, Radenković et al. 2011). Such a rich and diverse environment makes it particularly suitable for examining not only taxonomic but also functional diversity, as different types of habitat support different ecological functions (Gibb & Hochuli 2002). These habitats can be found in various vegetation types across SE Europe, and they display the dominant natural plant communities in accordance with current edaphic and climatic conditions (Bohn et al. 2007).

Based on data pertaining to selected functional traits of hoverflies in SE Europe, we aimed to
define hoverfly functional groups in this region based on the similarity of these traits, as well as
to compare the representation of delineated hoverfly functional groups among different
vegetation types.

78 MATERIAL AND METHODS

A list of all hoverflies in SE Europe and their representation in different vegetation types was
created based on data from the existing literature, personal observations (resulting from more
than 50 years of collecting hoverflies in the region; dataset deposited in the Database of
Department of Biology and Ecology, Faculty of Sciences, University of Novi Sad, Serbia), and
expert opinion.

Classification of hoverflies into functional groups was conducted based on 11 functional traits relating to the biological and ecological characteristics of each species, which together encompassed 46 trait categories. ... Information on larval microhabitat, larval feeding type, duration of larval development, inundation tolerance, number of generations and period of flight was directly extracted from existing literature (Speight 2018; Speight et al. 2020). Area of species range was inferred based on the available distribution data of analyzed species. Flight ability was partly based on field observations and partly on data available in Speight et al. 2020, from which information about the migratory status of species was extracted. Species was categorized as being a good or bad flier based on its ability to fly longer distances; species lingering around the feeding place, flying slowly and heavily were categorized as having bad flight ability, opposed to species crossing longer distances, flying fast and briskly, which were categorized as having good flight ability. Height at which species fly was scored solely on field observations. Species observed flying at height below 1m above the ground were classified as flying near the ground, while species flying above 1.5 m above the ground were scored as arboreal. Human impact tolerance was scored based on expert opinion. As data on body size was not available in the literature, we obtained these measures in the laboratory (details in Miličić et al. 2020). There were several possible states for each defined trait. To avoid bias for traits having multiple trait states, we applied a weighted average, whereby the weight of each trait state was divided by the number of trait states for that particular trait. In several cases where the trait state for a particular species was unknown, the species was assigned the most common trait state found among other species from the same genus. A list of the functional traits and corresponding trait states we considered is presented in Table S1.

As a preliminary analysis to categorize species into functional groups, we conducted a Principal Component Analysis (PCA). PCA was carried out by applying a normal varimax rotation of factor loadings (Dennis & Hellberg, 2010; Livshultz et al. 2011). Only factors with an eigenvalue greater than one were considered significant. Functional traits with a factor loading greater than 0.6 were considered relevant. To classify species into functional groups, we applied fuzzy

clustering in the R package cluster (Maechler et al. 2019). In fuzzy clustering, each observation can potentially belong to a larger number of clusters and thereby be "scattered" across a number of clusters (Podani 1994). To determine the optimal number of clusters, the Dunn partition coefficient was used, which varies between one for "hard" clusters and 1 / k (where k is the number of clusters) for fully dispersed datasets (Trauwaert 1988). We used silhouette width to determine the separation distance between clusters. Silhouette coefficients close to +1 indicate that a sample (species) is distant from neighboring clusters, a value of 0 indicates that a species is close to the boundaries of two clusters, and negative values indicate that a species is potentially included in the wrong cluster (Maechler et al. 2019). Allocation of species to one "hard cluster" was based on the highest percentage of attribution to some of the fuzzy clusters. After classification of the functional groups, the correlation of allocated variability with defined clusters was tested using analysis of variance (ANOVA). To examine the significance of the differences between clusters, a Tukey's HSD test was used. The degree to which individual species belonged to defined clusters (functional groups) was tested by Discriminant Function Analysis. Based on the square of the Mahalanobis distance, the UPGMA (Unweighted Pair-Group Method using Arithmetic Averages) method was applied to construct a dendrogram describing the distance between different functional groups.

In order to estimate the representation of functional groups of hoverflies in different vegetation types represented in SE Europe, we used the map of natural vegetation of Europe published in Miličić et al. (2020) (Fig. 1), which was based on a previously published map of natural vegetation in Europe (Bohn et al. 2000). Details on map conversion are reported in Miličić et al. (2020). A Chi-squared test was used to determine if there were significant differences between the frequencies of the different functional groups across 11 vegetation types in SE Europe.

RESULTS

In total, 568 species registered in SE Europe were included in the analyses. The first 5 PCA axes were kept in further analyses based on the results of the Scree test, explaining 39.9% of the variability (10.9%, 10.1%, 7.3%, 6.1% and 5.5%, respectively). The eigenvalues for axes 1-5 were 5.12, 4.77, 3.44, 2.85 and 2.60, respectively.

Division into functional groups

The fuzzy cluster analysis applied to the factor scores of first five PC axes resulted in five clearly separated functional groups (FGs) of hoverflies. Dunn's partitioning coefficient was 0.78 with a membership exponent set to 1.5. The average silhouette width for the total dataset was 0.52, with widths per cluster (FG) of 0.53, 0.29, 0.53, 0.62 and 0.70, respectively. Of the 568 analyzed species, 56 exhibited 100% affiliation to one FG, and a further 460 species were classified into an FG based on >70% affiliation. These parameters indicate that overall separation of species into FGs was good. There were 68 species in the first FG, 165 in the second, 78 in the third, 128 in the fourth, and 129 species in the fifth (Table S1).

The ANOVA showed that all five PCA axes describe differences among the defined FGs (Table S2). This outcome was further confirmed by the Tukey's HSD test, which identified significant differences (p < 0.05) and confirmed separation of the FGs based on all PCA axes (Table S3).

Table 1 shows functional traits of hoverflies that were significant for the separation of species into functional groups. FG1 consisted mainly of species with saprophagous larvae that develop in submerged sediment. These traits were negatively correlated with axis PC1, which clearly separated the first cluster (Fig. 2A). FG2 encloses widely distributed species (based on PC5, Fig. 2C) whose larvae are not tolerant to inundation, as defined by axes PC1 and PC4 (Fig. 2A). Axis PC2 can partly be used for the description of FG2, albeit cautiously, as it predominantly (but not exclusively) encompasses traits negatively correlated with PC2, designating species with very good flight ability that can migrate, have a high tolerance to human impact, and have more generations during the year to this group (Fig. 2B). However, FG1 consisted mainly of species with saprophagous larvae that develop in submerged sediment. These traits were negatively correlated with axis PC1, which clearly separated the first cluster (Fig. 2A). FG2 could be defined as comprising resilient and ubiquitous species, as this group encompasses species with good or very good flight ability that can migrate, have a high tolerance to human impact, and have more generations during the year. Axis PC2 revealed that variability among these latter traits is also reflected in FG1 (Fig. 2B) but, unlike FG1, FG2 includes species whose larvae are not tolerant to inundation, as defined by axis PC4. FG3 is clearly separated from other functional groups based on axis PC4 (Fig. 2C) It consisted mainly of species with saproxylic larvae, protracted larval development, and having less than one generation per year. FG4 was defined based on axis PC3 (Fig. 2B). It includes species whose larvae develop in plant roots, stems and leaves and with a low tolerance to human impact. FG5 consists of a high proportion of endemic and relict species with phytophagous larvae that develop in plant bulbs, as defined by axis PC5 (Fig. 2C). Additionally, axis PC4 revealed that the larvae of FG5 species are not tolerant to inundation (Table 1).

Discriminant analysis conducted on PC1-PC5 factor scores separated all hoverfly FGs with high significance based on the functional traits we considered (Table S4). Species were correctly classified into a priori-defined FGs with an overall accuracy of 92.79%: 97.06% of species within FG1, 92.73% in FG2, 98.72% in FG3, 88.29% in FG4, and 91.47% in FG5 were correctly classified.

The dendrogram based on the square of Mahalanobis distances revealed that species classified in
FG2 and FG4 exhibited the greatest similarity, whereas species in FG1 were the most distinct
based on the functional traits we used for classification (Fig. 3).

- 52 182 Functional groups and vegetation types
 53
- ⁵⁴ 183 Relative frequency of the different FGs varied significantly across SE Europe ($\chi^2(4) = 56.63$, p = 0.00), as well as across different vegetation types (Table 2). Species within FG2 dominated both

- - among SE European hoverflies and in the majority of vegetation types (9 of 11). The least
 represented species in the majority of vegetation types (6 of 11) were species within FG3,
 followed by FG1 and FG5, respectively (Table 2).

188 Interestingly, two vegetation types, south-west Balkan sub-Mediterranean mixed oak forests (H) 189 and Mediterranean mixed forests (J), did not follow this trend. With regard to the former, species 190 from FG4 were dominant ($\chi^2(4)$ = 50.64, p=0.00), whereas the latter contained the highest 191 percentage of species classified in FG5. South-west Balkan sub-Mediterranean mixed oak forests 192 are particularly interesting since they harbour the smallest percentages of species from both FG1 193 and FG3 (Fig. 4).

⁷ 194 **DISCUSSION**

19 195 Our multivariate classification of the functional traits of hoverflies identified five ecologically interpretable FGs. The significance of functional classification is that functional affiliation does not have to coincide with taxonomic similarities among species (Grime 1988). Thus, relationships between species can be revealed that could otherwise remain hidden if only a taxonomic classification is used.

- None of the defined FGs is genus-specific. Furthermore, although they dominated some of the FGs, the genera with the largest numbers of species (i.e., Cheilosia Meigen, 1838, Merodon Meigen, 1803, and *Eumerus* Meigen, 1822) were not exclusively grouped together in a single cluster. This arrangement of species within FGs confirms the notion that species may exhibit functional similarities, even though they exhibit significant differences in morphology and, moreover, that morphologically similar species may have different functions in the ecosystem (Young et al. 2007).
- Very few studies of the functional grouping of hoverflies have been conducted previously. Schweiger et al. (2007) analyzed 133 hoverfly species registered on agricultural land and the impact of intensive land use on the richness of functional groups. Keil et al. (2008) grouped 641 species of hoverflies recorded in Europe and then examined how richness of the groupings changed in relation to latitudinal variations, as well as the effect of selected environmental factors on functional richness. These studies have also revealed the importance of larval feeding type and larval inundation tolerance to delineating functional groups, though the relevance of some other traits differed from the results of our study. In both Schweiger et al. (2007) and Keil et al. (2008), traits that proved to be significant for functional categorization were larval microhabitat, number of generations per year, length of larval development, and body size. However, here, we also considered functional traits not assessed in those previous studies. Newly analyzed traits, such as flight ability and tolerance to human impact, had a great contribution to separating our functional groups. This outcome confirms that by considering a larger number of relevant characters, more comprehensive results can be obtained (Petchey & Gaston 2006).

Although our study was focused on hoverfly species in SE Europe, we hypothesize that this functional grouping could be applied to other regions as well. The reason lies in the fact that the traits that have proven to be most significant for the separation of the species into functional groups do not exhibit geographical variation, i.e. species would have the same state of a particular trait (e. g. larval feeding mode or extent of species range) in another region as well. Traits that vary across different geographical ranges, such as period of flight, were not marked as significant in defining hoverfly functional groups.

Most significant traits for the separation of hoverfly functional groups

The trait that proved dominant in distinguishing hoverfly FGs in our study is larval feeding type. Indeed, as for the extraordinary variability among adults, hoverfly larvae also display equally diverse feeding modes (Doyle et al. 2020). Considering the broad variation in this functional trait, which largely determines other biological and ecological characteristics of hoverfly species (Rotheray & Gilbert 2011), it is perhaps no surprise that we found this trait in particular to be the most significantly informative in terms of functional groups. For example, larval feeding type is directly related to the level of species specialization (Bonelli et al. 2011; Orsucci et al. 2018). Species having phytophagous larvae, which develop in roots and bulbs, are considered specialists, as are species with saproxylic larvae. This dependency on a host plant or, in the case of saproxylic species, a specific phase of tree decay has a considerable limiting effect on the possibility for a species to expand its range. Larvae of the saproxylic species *Blera fallax* (of FG3 herein) develop almost exclusively in rot holes of *Pinus sylvestris* (Rotheray et al. 2016). Species from the Cheilosia canicularis taxonomic group (which includes C. canicularis, C. hymantopus and C. ortotricha, and all classified within FG4) are even more specialized, with the larvae of these three species being exclusively associated with the plant *Petasites hybridus*, but each one develops in a different part of the plant, from root to leaf (Stuke & Claussen 2000).

We also found extent of species range to be an important parameter for defining functional groups, particularly FG5. Endemic or relict species are highly adapted to their particular niches (Harrison & Noss 2017), which limits their potential spread into new areas, as exemplified by the assignment to FG5 of many Merodon species restricted to specific Aegean islands (Radenković et al. 2011, Vujić et al. 2016).

Ability to fly determines the dispersal capability of hoverfly species; if a species is a good flyer it can migrate and expand the area of occupancy, whereas species that are poor flyers and only travel short distances have very limited distributions. Numerous studies have confirmed the significance of flight ability to resilience to extinction (Osborne et al. 2002; Chapman et al. 2015; Dällenbach et al. 2018, Chichorro et al. 2020).

Number of generations in a year can reflect the survival strategies of hoverfly species. Hoverflies that produce multiple generations annually usually produce large numbers of eggs (Zheng et al. 2019) and are considered less specific in terms of microhabitat selection (Speight et al. 2020).

However, species having fewer generations within a year are more likely to efficiently use
necessary resources compared to those having shorter generation timespans (Aguirre-Gutiérrez *et al.* 2016).

Inundation tolerance provides hoverflies with a superior survival kit upon exposure to wet conditions (Brust *et al.* 2007). Indeed, saprophagous hoverfly larvae that have breeding tubes enable them to survive in moist areas and more easily overcome challenging environmental conditions (Moquet *et al.* 2018).

Species tolerance to human impact has proved significant in defining the functional groups, revealing a link between the ability to resist changes in the environment caused by anthropogenic pressures and species functions in ecosystems (Samia *et al.* 2015).

Based on our results, the most functionally similar hoverfly species among those we considered are those of FG2 and FG4, with the most divergent being those in FG1. This similarity can be epitomized in the direct and indirect links between FG2, FG4 and FG5 and herbaceous plants. Larval development of species in FG4 and FG5 is related to different plant parts, and the hosts of larvae from FG2 species develop strictly within plant tissues, as many species from this group are aphidophagous. Species in FG3, characterized by having saproxylic larvae, are the second most divergent group. Unlike FG2, species in FG3 depend on dead plant matter, as these species are wood decomposers (Soszyńska-Maj et al. 2009). The reason why FG1 is the most divergent group is likely to the saprophagous nature of the larvae, which can be linked to extremely wet and, in many cases aquatic habitats, unlike for all other functional groups.

Functional groups and vegetation types

34 278

Species within FG4, dominant in south-west Balkan sub-Mediterranean mixed oak forests, mostly belong to the genus Cheilosia, whose larvae develop in roots, stems and leaves of host plants and that are particularly sensitive to anthropogenic impact. This high percentage of species directly dependent on specific vegetation types supports the notion that south-west Balkan sub-Mediterranean mixed oak forests represent an ecologically unique ecosystem, the high conservation value of which is often neglected (Mansourian et al. 2013). The diverse shrub understorev within these forests, intermixed with grasslands, increases habitat heterogeneity (Bugalho et al. 2011), supporting the macrohabitat requirements of these FG4 species. Notably, this vegetation type hosts the smallest percentage of species belonging to the functional groups FG1 and FG3, which mainly comprise saprophagous and saproxylic species, respectively. This pattern of FG representation may be attributable to climate change and inappropriate forest management. Indeed, climate change might have a particularly negative impact on species from FG1 in mixed oak forests, as these species are highly dependent on wet microhabitats (Speight 2018), which are severely affected by global warming (Papadopoulos & Pantera 2016). The second threat to forest health and condition is intensive forest management (whereby old oak trees and dead wood are removed from ecosystems), or even sometimes a lack of management

Entomological Science

(Stojanović *et al.* 2015). It is important to highlight the fact that this vegetation type exhibits the highest dark diversity of hoverflies in SE Europe (Miličić et al. 2020), which reflects reduced local biodiversity relative to potential richness. Therefore, it is likely that changes in management of such oak forests, such as retention of habitat trees (Mölder et al. 2020) and creating stepping-stones between veteran tree sites (Mestre et al. 2018), could potentially restore damaged ecosystems. In such circumstances, occurrence of a greater proportion of hoverflies dependent on dead or dying wood for some part of their lifecycle could be expected at such localities.

Mediterranean mixed forests were found to host a significant proportion of species within FG5, a group rich in endemic and relict species with phytophagous larvae that develop in bulbs. Mediterranean islands are characterized by high plant diversity and endemism (Georghiou & Delipetrou 2010), so a particular functional profile for species detected in Mediterranean mixed forests was somewhat anticipated. The long-lasting influence of human impacts in this region (Thompson 2005) has resulted in peculiar landscape patterns that have shaped distinctive species compositions. In particular, large and highly connected areas of this vegetation type, together with interspersed open habitats and the high diversity of bulbous plants (Petanidou et al. 2013), contribute to the maintenance of *Merodon* species that constitute a considerable proportion of the species in FG5.

CONCLUSION

We found that larval feeding type is the most dominant trait responsible for the categorization of 568 hoverfly species registered in SE Europe into five functional groups. The influence of different environmental pressures defines the richness of functional groups in specific vegetation types in this region. Further study is needed to investigate how established conservation measures can enable recovery of the functional richness affected by habitat disturbance, which would help us to further understand the importance of functional diversity to the preservation of biodiversity.

Acknowledgements: We kindly thank John O'Brien for English proofreading. This work was financially supported by the Ministry of Education, Science and Technological Development of the Republic of Serbia (Grant Nos. 451-03-9/2021-14/200358 and 451-03-68/2021-14/200125) and H2020 Project ANTARES, grant no. 664387.

References:

Page 11 of 26

1 2									
- 3 4 5	329 330	Aguirre-Gutiérrez J, Kissling WD, Carvalheiro LG, WallisDeVries MF, Franzén M, Biesmeijer JC (2016) Functional traits help to explain half-century long shifts in pollinator distributions.							
6	331	Scientific Reports 6, 1–13. 10.1038/srep24451							
7 8 0	332	Bohn U, Gollub G, Hettwer C, Weber H, Neuhäuslová Z, Raus T, Schlüter H (2000) Karte der							
9 10	333	natürlichen Vegetation Europas/Map of the Natural Vegetation of Europe. Maßstab/Scale 1:							
11 12	334	2,500,000.							
13	335	Bohn U, Zazanashvili N, Nakhutsrishvili G (2007) The Map of the Natural Vegetation of Europe							
14 15	336	and its application in the Caucasus Ecoregion. Bulletin of the Georgian National Academy of							
16	337	<i>Sciences</i> 175 , 112–121.							
17 18	338	Bonelli S. Witek M. Canterino S. Sielezniew M. Stankiewicz-Fiedurek A. Tartally A. Balletto F.							
19	339	Schönrogge K (2011) Distribution, host specificity and the potential for cryptic speciation in							
20 21	340	hoverfly <i>Microdon myrmicae</i> (Diptera: Syrphidae), a social parasite of <i>Myrmica</i> ants. <i>Ecological</i>							
22 23	341	<i>Entomology</i> 36 , 135–143. https://doi.org/10.1111/j.1365-2311.2010.01253.x							
24	342	Brust ML, Hoback WW, Wright RJ (2007) Immersion tolerance in rangeland grasshoppers							
25 26	343	(Orthoptera: Acrididae). Journal of Orthoptera Research 16, 135–138.							
27 28	344	https://doi.org/10.1665/1082-6467(2007)16[135:ITIRGO]2.0.CO;2							
29 20	345	Bugalho MN, Caldeira MC, Pereira JS, Aronson J, Pausas JG (2011) Mediterranean cork oak							
31	346	savannas require human use to sustain biodiversity and ecosystem services. Frontiers in Ecology							
32 33	347	and the Environment 9, 278–286. https://doi.org/10.1890/100084							
34 35	348	Cadotte MW, Carscadden K, Mirotchnick N (2011) Beyond species: functional diversity and the							
36	349	maintenance of ecological processes and services. Journal of Applied Ecology 48, 1079-1087.							
37 38	350	https://doi.org/10.1111/j.1365-2664.2011.02048.x							
39 40	351	Cardinale BJ, Duffy JE, Gonzalez A, Hooper DU, Perrings C, Venail P, Narwani A, Mace GM,							
40 41	352	Tilman D, Wardle DA, Kinzig AP (2012) Biodiversity loss and its impact on humanity. Nature							
42 43	353	486 , 59–67. 10.1038/nature11148							
44 45	354	Chapman JW, Reynolds DR, Wilson K (2015) Long-range seasonal migration in insects:							
45 46	355	mechanisms, evolutionary drivers and ecological consequences. Ecology Letters 18, 287-302.							
47 48	356	https://doi.org/10.1111/ele.12407							
49	357	Chichorro F, Urbano F, Teixeira D, Väre H, Pinto T, Brummitt N, He X, Hochkirch A, Hyvönen							
50 51	358	J, Kaila L, Juslén A, Cardoso P (2020) Species traits predict extinction risk across the Tree of							
52 53	359	Life. <i>BioRxiv</i> . doi: https://doi.org/10.1101/2020.07.01.183053							
54									
55 56									
57									
58		10							
59 60									

3 G0 Cornwell WK, Ackerly DD (2009) Community assembly and shifts in plant trait distributions across an environmental gradient in coastal California. *Ecological Monographs* 79, 109–126.
 3 6 https://doi.org/10.1890/07-1134.1

⁸ 363 Craven D, Filotas E, Angers, VA, Messier C (2016) Evaluating resilience of tree communities in
⁹ 364 fragmented landscapes: linking functional response diversity with landscape connectivity.
¹⁰ 365 *Diversity and Distributions* 22, 505–518. https://doi.org/10.1111/ddi.12423

 13
 14
 15
 16
 368
 Crnobrnja-Isailović J (2007) Cross-section of a refugium: genetic diversity of amphibian and reptile populations in the Balkans. In: Weiss S, Ferrand N (eds) *Phylogeography of Southern European Refugia*, pp 327–337. Springer, Dordrecht, The Netherlands.

¹⁸ 369 Dällenbach LJ, Glauser A, Lim KS, Chapman JW, Menz MH (2018) Higher flight activity in the
 ¹⁹ 370 offspring of migrants compared to residents in a migratory insect. *Proceedings of the Royal* ²¹ 371 Society B 285, 20172829. 10.1098/rspb.2017.2829

- ²³ 372 Davis ES, Kelly R, Maggs CA, Stout JC (2018) Contrasting impacts of highly invasive plant
 ²⁴ 373 species on flower-visiting insect communities. *Biodiversity and Conservation* 27, 2069–2085.
 ²⁶ 374 10.1007/s10531-018-1525-y
- ²⁸ 375 Day RL, Hickman JM, Sprague RI, Wratten SD (2015) Predatory hoverflies increase oviposition
 ²⁹ 376 in response to colour stimuli offering no reward: implications for biological control. *Basic and* ³¹ 377 Applied Ecology 16, 544–552. 10.1016/j.baae.2015.05.004
- 378 Dennis AB, Hellberg ME (2010) Ecological partitioning among parapatric cryptic
 379 species. *Molecular ecology*, 19, 3206–3225.
- ³⁶
 ³⁷
 ³⁸
 ³⁸
 ³⁸
 ³⁹
 ³⁸
 ³⁹
 <li
- ⁴¹
 ⁴² 383 Doyle T, Hawkes WL, Massy R, Powney GD, Menz MH, Wotton KR (2020) Pollination by
 ⁴³ 384 hoverflies in the Anthropocene. *Proceedings of the Royal Society B* 287, 20200508.
 ⁴⁴ 385 https://doi.org/10.1098/rspb.2020.0508
- Georghiou K, Delipetrou P (2010) Patterns and traits of the endemic plants of Greece. *Botanical Journal of the Linnean Society* 162, 130–153. https://doi.org/10.1111/j.1095-8339.2010.01025.x
- Gibb H, Hochuli DF (2002) Habitat fragmentation in an urban environment: large and small
 fragments support different arthropod assemblages. *Biological Conservation* 106, 91–100.
 https://doi.org/10.1016/S0006-3207(01)00232-4

11

57 58

55 56

1 2

7

12

17

22

27

32

Page 13 of 26

1 2		
3	391	Gitay H, Noble IR (1997) What are functional types and how should we seek them? In: Smith
4 5	392	TM, Shugart HH, Woodward FI (eds) Plant functional types: their relevance to ecosystem
6 7	393	properties and global change, pp 3-19. Cambridge University Press, Cambridge.
8	394	Grime JP (1988) The CSR model of primary plant strategies—origins, implications and tests. In:
9 10 11	395	Gottlieb L, Jain S (eds) Plant Evolutionary Biology, pp 371–393. Springer Netherlands.
12	396	Grosskopf G (2005) Biology and life history of Cheilosia urbana (Meigen) and Cheilosia
13	397	psilophthalma (Becker), two sympatric hoverflies approved for the biological control of
14 15	398	hawkweeds (Hieracium spp.) in New Zealand. Biological Control 35, 142-154.
16	399	https://doi.org/10.1016/j.biocontrol.2005.06.013
17		
18	400	Harrison S, Noss R (2017) Endemism hotspots are linked to stable climatic refugia. Annals of
19 20 21	401	Botany 119, 207–214. https://doi.org/10.1093/aob/mcw248
22	402	Hooper D, Buchmann N, Degrange V, Díaz SM, Gessner M, Grime P, Hulot F, Mermillod-
23	403	Blondin F, van Peer L, Roy J, Symstad A, Solan M, Spehn EM (2002) Species diversity,
24	404	functional diversity and ecosystem functioning. In: Loreau M. Naeem S. Inchausti P (eds)
25 26	405	<i>Biodiversity and Ecosystem Functioning</i> pp 195–208 Oxford University Press Oxford
27		
28	406	Jauker F, Wolters V (2008) Hover flies are efficient pollinators of oilseed rape. Oecologia 156,
29 30	407	819–823. 10.1007/s00442-008-1034-x
31 32	408	Keil P. Dziock F. Storch D (2008) Geographical patterns of hoverfly (Diptera, Syrphidae)
33	409	functional groups in Europe inconsistency in environmental correlates and latitudinal trends
34 35	410	<i>Ecological Entomology</i> 33 , 748–757. https://doi.org/10.1111/j.1365-2311.2008.01032.x
36	411	Li P. Kleijn D. Badenhausser I. Zaragoza-Trello C. Gross N. Raemakers I. Schener I (2020) The
37 38	/12	relative importance of green infrastructure as refuge habitat for pollinators increases with local
39	712 //12	land use intensity. <i>Journal of Applied Ecology</i> 57, 1494, 1503, https://doi.org/10.1111/1365
40	415	2664 12659
41 42	414	2004.13038
42 43	415	Livshultz T, Mead JV, Goyder DJ, Brannin M. (2011) Climate niches of milkweeds with
44	416	plesiomorphic traits (Secamonoideae: Apocynaceae) and the milkweed sister group link ancient
45 46	417	African climates and floral evolution <i>American Journal of Botany</i> 98 1966–1977
40 47	,	
48	418	Maechler M, Rousseeuw P, Struyf A, Hubert M, Hornik K (2019) cluster: Cluster Analysis
49	419	Basics and Extensions. R package version 2.1.0. Available from URL: https://cran.r-
50	420	project org/web/packages/cluster/index html
51 52	.20	project.org, web, pachages, craster, machinin
53	421	Mansourian S, Rossi M, Vallauri D (2013) Ancient forests in the northern Mediterranean:
54	422	neglected high conservation value areas. Marseille, WWF France. 80p.
55		6
56 57		
58		12
59		
60		

3 Medellín RA, Equihua M, Amin MA (2000) Bat diversity and abundance as indicators of 423 4 Neotropical rainforests. biology 1666-1675. 424 disturbance in Conservation 14. 5 https://doi.org/10.1111/j.1523-1739.2000.99068.x 6 425

Mestre L, Jansson N, Ranius T (2018) Saproxylic biodiversity and decomposition rate decrease
with small-scale isolation of tree hollows. *Biological Conservation* 227, 226–232.
https://doi.org/10.1016/j.biocon.2018.09.023

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.
 https://doi.org/10.1007/s10531-017-1486-6

Miličić M, Popov S, Vujić A, Ivošević B, Cardoso P (2020) Come to the dark side! The role of functional traits in shaping dark diversity patterns of south-eastern European hoverflies.
 Ecological Entomology 45, 232–242. https://doi.org/10.1111/een.12788

435 Moquet L, Laurent E, Bacchetta R, Jacquemart AL (2018) Conservation of hoverflies (Diptera,
436 Syrphidae) requires complementary resources at the landscape and local scales. *Insect*437 *Conservation and Diversity* 11, 72–87. https://doi.org/10.1111/icad.12245

438 Mölder A, Schmidt M, Plieninger T, Meyer P (2020) Habitat-tree protection concepts over 200
439 years. *Conservation Biology* 34, 1444–1451. https://doi.org/10.1111/cobi.13511

440 Ooi MK, Denham AJ, Santana VM, Auld TD (2014) Temperature thresholds of physically
 441 dormant seeds and plant functional response to fire: variation among species and relative impact
 442 of climate change. *Ecology and Evolution* 4, 656–671. https://doi.org/10.1002/ece3.973

Orsucci M, Audiot P, Dorkeld F, Pommier A, Vabre M, Gschloessl B, Rialle S, Severac D,
 Bourguet D, Streiff R (2018) Larval transcriptomic response to host plants in two related
 phytophagous lepidopteran species: implications for host specialization and species divergence.
 BMC genomics 19, 265. https://doi.org/10.1186/s12864-018-4589-x

43 447 Osborne JL, Loxdale HD, Woiwod IP (2002) Monitoring insect dispersal: methods and
44 448 approaches. In: Bullock JM, Kenward RE, Hails RS (eds) *Dispersal Ecology: 42nd Symposium*449 of the British Ecological Society, pp 24–49. Cambridge University Press, Cambridge.

48 450 Papadopoulos A, Pantera A (2016) Dendrochronological Investigations of Valonia Oak Trees in
 49 451 Western Greece. South-east European forestry 7, 29–37. https://doi.org/10.15177/seefor.16-05

452 Perkins MJ, Inger R, Bearhop S, Sanders D (2018) Multichannel feeding by spider functional
 453 groups is driven by feeding strategies and resource availability. *Oikos* 127, 23–33.
 454 https://doi.org/10.1111/oik.04500

13

56 57

42

47

1 2

7

12

17

22

27

58

Page 15 of 26

1 2								
3	455	Persson AS, Ekroos J, Olsson P, Smith HG (2020) Wild bees and hoverflies respond differently						
4	456	to urbanisation, human population density and urban form. <i>Landscape and Urban Planning</i> 204 ,						
5 6 7	457	103901. https://doi.org/10.1016/j.landurbplan.2020.103901						
8	458	Petanidou T, Ståhls G, Vujić A, Olesen JM, Rojo S, Thrasyvoulou A, Sgardelis S, Kallimanis						
9 10	459	AS, Kokkini S, Tscheulin T (2013) Investigating plant—pollinator relationships in the Aegean:						
10	460	the approaches of the project POL-AEGIS (The pollinators of the Aegean archipelago: diversity						
12	461	and threats) Journal of Anicultural Research 52 106–117						
13	462	https://doi.org/10.3896/IBRA 1.52.2.20						
14 15	102	https://doi.org/10.3090/10101111.1.32.2.20						
16	463	Petchey OL, Gaston KJ (2006) Functional diversity: back to basics and looking forward. Ecology						
17 18	464	<i>Letters</i> 9, 741–758. https://doi.org/10.1111/j.1461-0248.2006.00924.x						
19	465	Peters VE, Campbell KU, Dienno G, García M, Leak E, Lovke C, Ogle M, Steinly B, Crist, TO						
20 21	466	(2016) Ants and plants as indicators of biodiversity ecosystem services and conservation value						
22	167	in constructed grasslands <i>Riodiversity</i> and <i>Conservation</i> 25 1481–1501						
23	467	https://doi.org/10.1007/s10531_016_1120_7						
24	400	https://doi.org/10.100//310551-010-1120-2						
25 26	469	Podani J (1994) Multivariate Data Analysis in Ecology and Systematics Ecological						
27	470	Computations Series (ECS) Vol 6 SPB Academic Publishing The Hague The Netherlands						
28	470	computations series (Leo), vol. 0. Si D ricadenne i donsning, The Hagae, The Realerando.						
29	471	Popov S, Miličić M, Diti I, Marko O, Sommaggio D, Markov Z, Vujić A (2017) Phytophagous						
30 31	472	hoverflies (Diptera: Syrphidae) as indicators of changing landscapes Community Ecology 18						
32	473	287_294 https://doi.org/10.1556/168.2017.18.3.7						
33	175							
34 25	474	Poulakakis N, Kapli P, Lymberakis P, Trichas A, Vardinoyiannis K, Sfenthourakis S, Mylonas						
35 36	475	M (2015) A review of phylogeographic analyses of animal taxa from the Aegean and						
37	476	surrounding regions. Journal of Zoological Systematics and Evolutionary Research 53, 18–32.						
38	477	https://doi.org/10.1111/izs.12071						
39 40								
40 41	478	Radenković S, Vujić A, Ståhls G, Perez-Banon C, Rojo S, Petanidou T, Šimić S (2011) Three						
42	479	new cryptic species of the genus Merodon Meigen (Diptera: Syrphidae) from the island of						
43	480	Lesvos (Greece). Zootaxa 2735, 35–56.						
44 45								
45 46	481	Rotheray GE, Gilbert F (2011) The natural history of hoverflies. Ceredigion, Forrest text.						
47								
48	482	Rotheray EL, Goulson D, Bussiere LF (2016) Growth, development, and life-history strategies in						
49 50	483	an unpredictable environment: case study of a rare hoverfly Blera fallax (Diptera, Syrphidae).						
50 51	484	Ecological Entomology 41, 85–95. https://doi.org/10.1111/een.12269						
52								
53	485	Sabovljević M, Natcheva R, Dihoru G, Tsakiri E, Dragićević S, Erdağ A, Papp B (2008) Check-						
54 57	486	list of the mosses of SE Europe. Phytologia Balcanica 14, 207–244.						
55 56								
57								
58		14						
59 60								
00								

Samia DS, Nakagawa S, Nomura F, Rangel TF, Blumstein DT (2015) Increased tolerance to

- wildlife. Nature *Communications* humans among disturbed 6, 1 - 8. https://doi.org/10.1038/ncomms9877 Schweiger O, Musche M, Bailey D, Billeter R, Diekötter T, Hendrickx F, Herzog F, Liira J, Maelfait JP, Speelmans M, Dziock F (2007) Functional richness of local hoverfly communities (Diptera, Syrphidae) in response to land use across temperate Europe. Oikos 116, 461-472. https://doi.org/10.1111/j.2007.0030-1299.15372.x Soszyńska-Maj A, Soszyński B, Klasa A (2009) Distribution and ecology of the saproxylic hoverfly Chalcosyrphus eunotus (Loew, 1873) (Diptera: Syrphidae) in Poland. Fragmenta Faunistica 52, 191–195. Speight MCD (2018) Species accounts of European Syrphidae, 2018. In: Speight MCD, Castella E, Sarthou JP, Vanappelghem C (eds) Syrph the Net, the Database of European Syrphidae (Diptera), Vol. 103, p 302. Syrph the Net publications, Dublin. Speight MCD, Castella E, Sarthou JP (2020) Syrph the Net 2020. In: Speight MCD, Castella E, Sarthou JP, Vanappelghem C (eds) Syrph the Net on CD, Issue 12, Syrph the Net Publications, Dublin. Stojanović DB, Levanič T, Matović B, Orlović S (2015) Growth decrease and mortality of oak floodplain forests as a response to change of water regime and climate. European Journal of Forest Research 134, 555–567. https://doi.org/10.1007/s10342-015-0871-5 Stuke JH, Claussen C (2000) Cheilosia canicularis auctt.- a complex of species. Volucella 5, 79-94. Szigeti V, Fenesi A, Soltész Z, Berki B, Kovács-Hostyánszki A (2020) Neutral effect of an invasive plant species with specialized flower structure on native pollinator communities. Biological Invasions 22, 3017-3030. https://doi.org/10.1007/s10530-020-02305-6 Thompson JN (2005) Coevolution: the geographic mosaic of coevolutionary arms races. Current Biology 15, R992-R994. https://doi.org/10.1016/j.cub.2005.11.046 Trauwaert E (1988) On the meaning of Dunn's partition coefficient for fuzzy clusters. Fuzzy sets and systems 25, 217-242. Vujić A, Petanidou T, Tscheulin T, Cardoso P, Radenković S, Stahls G, Baturan Ž, Mijatović G, Rojo S, Perez-Banon C, Devalez J, Andrić A, Jovičić S, Krašić D, Markov Z, Radišić D, Tataris G (2016) Biogeographical patterns of the genus Merodon Meigen, 1803 (Diptera: Syrphidae) in islands of the eastern Mediterranean and adjacent mainland. Insect Conservation and Diversity 9, 181-191. https://doi.org/10.1111/icad.12156

1 2 3 4 5	520 521	Vujić A, Šimić S, Radenković S (2001) Endangered species of hoverflies (Diptera: Syrphidae) on the Balkan Peninsula. <i>Acta Entomologica Serbica</i> 5 , 93–105.
6 7 8 9 10	522 523 524	Winsa M, Öckinger E, Bommarco R, Lindborg R, Roberts SP, Wärnsberg J, Bartomeus I (2017) Sustained functional composition of pollinators in restored pastures despite slow functional restoration of plants. <i>Ecology and Evolution</i> 7 , 3836–3846. https://doi.org/10.1002/ece3.2924
11 12 13 14 15	525 526 527	Young RL, Haselkorn TS, Badyaev AV (2007) Functional equivalence of morphologies enables morphological and ecological diversity. <i>Evolution</i> 61 , 2480–2492. https://doi.org/10.1111/j.1558-5646.2007.00210.x
15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43	528 529 530	Zheng Z, Liu H, Wang X, Wu X, Chen Y, Deng J, Chen X, Yuejian L, Pu D (2019) Development and reproduction of the hoverfly <i>Eupeodes corollae</i> (Diptera: Syrphidae). <i>Journal</i> of Earth Sciences & Environmental Studies 4, 4.
44 45 46 47 48 49 50 51		

Figure legends:

Figure 1. Map of vegetation types in Southeast Europe. A - Alpine; subalpine and oro-Mediterranean vegetation; B - montane spruce and mixed spruce forests; C - montane pine forests; D - acidophilous oak and mixed oak-hornbeam forests; E - beech and mixed beech forests; F - thermophilous mixed bitter, pedunculate or sessile oak forests; G - south-east Balkan sub-Mediterranean mixed oak forests; H - south-west Balkan sub-Mediterranean mixed oak forests; I - Pannonian lowland mixed oak forests and steppes; J - Mediterranean mixed forests; K - hardwood alluvial forests, wet lowland forests and swamps. Map is published in Miličić *et al.* (2020), available at: https://onlinelibrary.wiley.com/doi/full/10.1111/een.12788.

Figure 2. The distribution of clusters of functional groups, as defined by PCA axes. a) Axis PC1 clearly separates FG1 from other functional groups, whereas axis PC4 differentiates FG2 and FG3. b) Axis PC2 displays the variability between FG1 and FG2, whereas axis PC3 differentiates FG2 and FG4. c) Axis PC4 clearly separates FG3 from other functional groups, and axis PC5 differentiates FG5.

Figure 3. The UPGMA dendrogram constructed based on the square of the Mahalanobis distances depicts the similarity among the defined functional groups.

Figure 4. A comparison of the number of hoverfly species present in the different functional groups across 11 vegetation types in Southeast Europe. A - Alpine; subalpine and oro-Mediterranean vegetation; B - montane spruce and mixed spruce forests; C - montane pine forests; D - acidophilous oak and mixed oak-hornbeam forests; E - beech and mixed beech forests; F - thermophilous mixed bitter, pedunculate or sessile oak forests; G - south-east Balkan sub-Mediterranean mixed oak forests; H - south-west Balkan sub-Mediterranean mixed oak forests; I - Pannonian lowland mixed oak forests and steppes; J - Mediterranean mixed forests; K - hardwood alluvial forests, wet lowland forests and swamps.

 Table legends: Table 1. Results of PCA analysis of traits used for functional classification of hoverflies. Factor loading values greater than ± 0.6 are bolded and underlined.

Table 2. Functional group composition (relative frequency percentage of hoverfly species) of the hoverfly community across different vegetation types in South East Europe. A - Alpine; subalpine and oro-Mediterranean vegetation; B - montane spruce and mixed spruce forests; C - montane pine forests; D - acidophilous oak and mixed oak-hornbeam forests; E - beech and mixed beech forests; F - thermophilous mixed bitter, pedunculate or sessile oak forests; G - south-east Balkan sub-Mediterranean mixed oak forests; H - south-west Balkan sub-Mediterranean mixed oak forests; J - Mediterranean mixed forests; K - hardwood alluvial forests, wet lowland forests and swamps.

SUPPORTING INFORMATION

Additional Supporting Information may be found online in the Supporting Information section at the end of the article.

Table S1. List of all analyzed hoverfly species with its attribution to functional groups (FGs), percentage of belonging to each of the clusters and silhouette width.

Table S2. Analysis of variance of defined functional groups.

Table S3. Results of Tuckey HSD test for PC1-PC5.

Table S4. Statistical significance of the difference between defined functional groups based on discriminant analysis. p values - below the diagonal; F values - above the diagonal; df = 5.56



Figure 1. Map of vegetation types in Southeast Europe. A - Alpine; subalpine and oro-Mediterranean vegetation; B - montane spruce and mixed spruce forests; C - montane pine forests; D - acidophilous oak and mixed oak-hornbeam forests; E - beech and mixed beech forests; F - thermophilous mixed bitter, pedunculate or sessile oak forests; G - south-east Balkan sub-Mediterranean mixed oak forests; H - south-west Balkan sub-Mediterranean mixed oak forests; J - Mediterranean mixed forests; K - hardwood alluvial forests, wet lowland forests and swamps. Map is published in Miličić et al. (2020), available at: https://onlinelibrary.wiley.com/doi/full/10.1111/een.12788.

297x210mm (598 x 598 DPI)





Figure 2. The distribution of clusters of functional groups, as defined by PCA axes. A) Axis PC1 clearly separates FG1 from other functional groups, whereas axis PC4 differentiates FG2 and FG3. B) Axis PC2 displays the variability between FG1 and FG2, whereas axis PC3 differentiates FG2 and FG4. C) Axis PC4 clearly separates FG3 from other functional groups, and axis PC5 differentiates FG5.

177x371mm (96 x 96 DPI)



similarity among the defined functional groups.

166x109mm (96 x 96 DPI)



Figure 4. A comparison of the number of hoverfly species present in the different functional groups across 11 vegetation types in Southeast Europe. A - Alpine; subalpine and oro-Mediterranean vegetation; B - montane spruce and mixed spruce forests; C - montane pine forests; D - acidophilous oak and mixed oak-hornbeam forests; E - beech and mixed beech forests; F - thermophilous mixed bitter, pedunculate or sessile oak forests; G - south-east Balkan sub-Mediterranean mixed oak forests; H - south-west Balkan sub-Mediterranean mixed oak forests; J - Mediterranean mixed forests; K - hardwood alluvial forests, wet lowland forests and swamps.

196x133mm (149 x 149 DPI)

Table 1. Results of PCA analysis of traits used for functional classification of hoverflies. Factor loading values greater than ± 0.6 are bolded and underlined.

Trait	Trait state	PC1	PC2	PC3	PC4	PC5
Larval microhabitat	trees	0.18	-0.10	0.37	-0.48	0.38
	upward climbing lianas	0.07	-0.38	0.12	0.03	0.04
	herb layer	0.43	-0.01	-0.31	0.39	-0.20
	timber	0.02	0.05	-0.11	-0.57	0.03
	dung	-0.28	-0.23	-0.02	0.00	0.01
	litter	0.02	-0.09	0.05	0.02	0.13
	stones	0.01	0.03	0.03	-0.01	0.05
	nests of social insects	0.04	0.05	0.15	0.10	0.08
	root zone	0.28	0.29	-0.21	0.26	-0.38
	water plants	-0.45	0.07	0.14	0.08	0.05
	submerged sediment/debris	<u>-0.87</u>	-0.05	-0.04	0.05	0.01
Larval feeding mode	saprophagous	<u>-0.94</u>	-0.08	0.01	0.08	0.02
	saproxylic	-0.01	0.06	0.03	<u>-0.89</u>	0.08
	phytophagous-bulbs	0.19	0.14	0.09	0.19	<u>-0.74</u>
	phytophagous-roots	0.16	0.17	<u>-0.68</u>	0.22	0.14
	zoophagous	0.36	-0.26	0.42	0.29	0.46
Duration of larval development	less than 2 months	-0.08	-0.59	0.04	0.03	0.08
	2-6 months	-0.08	-0.12	0.02	0.18	0.02
	7-12 months	0.04	0.26	-0.08	0.28	-0.02
	more than a year	0.10	0.03	0.07	<u>-0.77</u>	-0.03
Inundation tolerance	intolerant	<u>0.65</u>	-0.07	0.03	<u>0.61</u>	-0.15
	tolerant (short breathing tube)	-0.01	0.16	-0.1	-0.09	0.16
	tolerant (medium breathing	0.4	0.00	0.02	0.40	0.00
	lube)	-0.4	0.09	-0.02	-0.49	0.08
Number of generations	loss then one consistion	-0.52	-0.13	0.08	-0.28	0.00
Number of generations	less than one generation	0.08	0.03	0.10	<u>-0.07</u>	-0.02
		0.03	0.23	-0.07	0.08	-0.11
	two generations	-0.02	0.05	0.03	0.17	0.1
Devied of flight		-0.1	-0.64	0.00	0.07	0.06
Period of flight	early spring	0.03	-0.14	0.01	0.03	-0.05
	spring	0.02	0.07	-0.07	-0.03	-0.02
	early summer	-0.06	0.06	-0.18	-0.06	0.28
	summer	0.01	0.02	0.09	0.05	0.05
		0.01	-0.09	0.16	0.02	-0.27
Boay size	small	0.03	0.01	-0.05	0.06	0.24
	medium	0.01	-0.07	0.01	0.14	-0.24
	large	-0.05	0.07	0.05	-0.23	0.05
Area of species range	endemic and/or relict	0.06	0.03	-0.02	-0.01	<u>-0.82</u>
	widely distributed	-0.06	-0.03	0.02	0.01	<u>0.82</u>

Flight ability	very good-migrants	0.03	<u>-0.76</u>	-0.04	0.03	0.04
	good	-0.04	<u>0.70</u>	0.05	0.00	0.00
	bad	0.04	-0.07	-0.01	-0.03	-0.05
Height at which species fly	arboreal	0.12	0.00	-0.23	-0.19	0.43
	near the ground	-0.12	0.00	0.23	0.19	-0.43
	low	0.11	0.10	<u>-0.79</u>	-0.02	-0.03
Human impact tolerance	medium	0.04	0.26	<u>0.82</u>	-0.03	-0.01
	high	-0.24	-0.08	-0.07	0.11	0.07
	very high	-0.10	<u>-0.8</u>	-0.08	0.01	0.02

to Review Only

Table 2. Functional group composition of the hoverfly community across different vegetation types in South East Europe. A - Alpine; subalpine and oro-Mediterranean vegetation; B - montane spruce and mixed spruce forests; C -montane pine forests; D - acidophilous oak and mixed oak-hornbeam forests; E - beech and mixed beech forests; F - thermophilous mixed bitter, pedunculate or sessile oak forests; G - south-east Balkan sub-Mediterranean mixed oak forests; H - south-west Balkan sub-Mediterranean mixed oak forests; I - Pannonian lowland mixed oak forests and steppes; J - Mediterranean mixed forests; K - hardwood alluvial forests, wet lowland forests and swamps; No. species - number of species per functional group.

Vege	etation type	FG1	FG2	FG3	FG4	FG5	Total	χ^2 value
А	No. species	20	79	16	71	24	210	
	% within A	9.52	37.62	7.62	33.81	11.43	100	$\chi^{2}(4) = 56.62/55,$ p= 00000
	% within total sample	3.52	13.91	2.82	12.5	4.23	36.97	P
В	No. species	28	98	18	86	19	249	2(4) 102 0100
	% within B	11.24	39.36	7.23	34.54	7.63	100	$\chi^{2}(4) = 102.8108$, p=.00000
	% within total sample	4.93	17.25	3.17	15.14	3.35	43.84	P
С	No. species	26	72	23	47	36	204	2(4) 0 4(2 420
	% within C	12.75	35.29	11.27	23.04	17.65	100	$\chi^{2}(4) = 9.463429,$ p=.05050
	% within total sample	4.58	12.68	4.05	8.27	6.34	35.92	P
D	No. species	24	86	27	65	24	226	2(4) 42 5207(
	% within D	10.62	38.05	11.95	28.76	10.62	100	$\chi^{2}(4) = 42.53976,$ p=.00000
	% within total sample	4.23	15.14	4.75	11.44	4.23	39.79	P
Е	No. species	51	136	56	104	53	400	2(4) 72 (7274
	% within E	12.75	34.00	14.00	26.00	13.25	100	$\chi^{2}(4) = 72.67274,$ p=.00000
	% within total sample	8.98	23.94	9.86	18.31	9.33	70.42	P
F	No. species	32	91	26	66	42	257	2(4) 01 52501
	% within F	12.45	35.41	10.12	25.68	16.34	100	$\chi^{2}(4) = 21.53/01,$ p=.00025
	% within total sample	5.63	16.02	4.58	11.62	7.39	45.24	r
G	No. species	38	90	42	62	48	280	2(4) 11 2227(
	% within G	13.57	32.14	15.00	22.14	17.14	100	$\chi^{2}(4) = 11.22276,$ p=.02417
	% within total sample	6.69	15.85	7.39	10.92	8.45	49.3	I ··· ·
Н	No. species	9	26	6	55	40	136	2(4) 50 (4192
	% within H	6.62	19.12	4.41	40.44	29.41	100	$\chi^{-}(4) = 50.64183,$ p=.00000
	% within total sample	1.58	4.58	1.06	9.68	7.04	23.94	r
Ι	No. species	41	73	22	50	19	205	-2(4) = 50,12945
	% within I	20.00	35.61	10.73	24.39	9.27	100	$\chi^{2}(4) = 50.13845,$ p=.00000
	% within total sample	7.22	12.85	3.87	8.8	3.35	36.09	r
J	No. species	31	68	42	31	102	274	2(4) 02 100((
	% within J	11.31	24.82	15.33	11.31	37.23	100	$\chi^{2}(4) = 83.12266,$ p=.00000
	% within total sample	5.46	11.97	7.39	5.46	17.96	48.24	1
Κ	No. species	37	74	27	55	29	222	2(4) 05 41514
	% within K	16.67	33.33	12.16	24.77	13.06	100	$\chi^{-}(4) = 25.41516,$ p=.00004
	% within total sample	6.51	13.03	4.75	9.68	5.11	39.08	