Distribution patterns of Hydropsychids and Rhyacophilids species (Trichoptera) in a not regulated Mediterranean river (SW Spain)

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ABSTRACT

Distribution patterns of Hydropsychids and Rhyacophilids species (Trichoptera) in a non-regulated Mediterranean river (SW Spain)

This paper investigates the longitudinal ordination of the Hydropsychidae and Rhyacophilidae species present in the high basin of the Hozgarganta River. The ordination of the Hydropsychids species present in the basin show three distribution patterns: Diplectrona felix and Hydropsyche infernalis are confined to the head streams; H. siltalai and H. iberomaroccana are distributed throughout the entire study zone, although the second one does not reach the highest sections of the basin; finally, H. lobata occupies the lowest sections, specially in the main axis of the river. The two Rhyacophila species studied show also a well differentiated distribution. R. fonticola is more abundant in the gorges and high sections, whereas R. munda prefers the riverbed of the main river, although it’s also found in some tributaries. Even in these intermediate sections, the segregation between the two species is almost perfect.

The water permanence in the riverbeds influences the distribution of some species. We have found that D. felix and H. infernalis inhabit the permanent sections of the stream heads whereas H. iberomaroccana significantly dominates the intermittent ones. H. siltalai, H. lobata, R. fonticola, and R. munda do not show a significant preference for any type of riverbed (permanent, intermittent, or ephemeral). It is interesting to highlight the survival of some H. iberomaroccana larvae in isolated pools during the summer. We suggest the possibility that these larvae survive thanks to the existence of a weak underground flow of subsurface origin between adjacent pools.

Key words: Hydropsychidae, Rhyacophilidae, distribution, seasonality, Hozgarganta river basin.

RESUMEN

Patrones de distribución de especies de Hydropsychidae y Rhyacophilidae (Trichoptera) en un río mediterráneo no regulado (Suroeste de la Península Ibérica)

En el presente trabajo se investiga la ordenación longitudinal de las especies de Hydropsychidae y Rhyacophilidae presentes en la cuenca alta del río Hozgarganta. La ordenación de las especies de Hydropsychidae presentes en la cuenca muestra tres patrones de distribución: Diplectrona felix e Hydropsyche infernalis están confinadas en los arroyos de cabecera; H. siltalai e H. iberomaroccana están distribuidas por toda la zona de estudio, aunque la segunda no alcanza los tramos más altos de la cuenca; por último, H. lobata ocupa los tramos más bajos, especialmente en el eje principal del río. Las dos especies de Rhyacophila estudiadas también muestran una distribución bien diferenciada. R. fonticola es más abundante en las gargantas y tramos altos, mientras que R. munda prefiere el cauce del río principal, aunque también se encuentra presente en algunos tributarios. Incluso en estos tramos intermedios la segregación entre ambas especies es casi perfecta.

La permanencia del agua en los cauces influye en la distribución de algunas especies. Hemos encontrado que D. felix e H. infernalis habitan los tramos permanentes de cabecera, mientras que H. iberomaroccana domina de forma significativa en los intermitentes. H. siltalai, H. lobata, R. fonticola y R. munda no muestran una preferencia significativa por algún tipo de cauce (permanent, intermitente o efímero). Es interesante resaltar la supervivencia de algunas larvas de H. iberomaroccana en pozas aisladas durante el verano. Sugerimos la posibilidad de que estas larvas sobrevivan gracias a la existencia de un débil flujo subterráneo de origen freático entre pozas adyacentes.

Palabras clave: Hydropsychidae, Rhyacophilidae, distribución, estacionalidad, cuenca río Hozgarganta.
INTRODUCTION

Both Hydropsychids and Rhyacophilids are two of the most diverse families of caddisflies in the Iberian Peninsula, in addition to the Hydroptilidae (González et al., 1987) and they are widespread in all Spanish catchments.

The ecological differences between both families are a consequence of the way of using the silk (Mackay & Wiggins, 1979). The Rhyacophilids have free-living larvae that only use the silk before pupation, whereas the Hydropsychidae are sedentary and net-building along with other filter-feeders, therefore they perform an important function in the treatment of the organic matter in fluvial ecosystems (Basaguren & Orive, 1991). In the Mediterranean rivers they are, together with the Hydroptilidae, the most important family of caddisflies due to their abundance and species’ richness (Giudicelli et al., 1985).

Many studies have pointed out a longitudinal replacement of Hydropsychidae species (García de Jalón, 1986, Camargo, 1992, Gallardo-Mayenco et al., 1998). The environmental factors change along the watercourse and the species occupying a particular position in the longitudinal sequence seem to have appropriate suites of physiological and behavioural characteristics (Edington & Hildrew, 1995).

Mediterranean climatic regions are distributed around the world, and seasonality and variability in rainfall is its main characteristic. Although the seasonal precipitation pattern is highly predictable in Mediterranean climatic areas, annual rainfall can vary largely from year to year (Gasith & Resh, 1999). As a consequence of this climatic pattern, flow irregularity is one of the most important features of the Mediterranean rivers (Giudicelli et al., 1985), with a seasonal pattern of high discharge during the wet period, which is followed by low discharge in the summer. Although, in some areas, they can be dry.

Aquatic insects that live in temporary ponds have developed a series of strategies that allow them to survive a periodic loss of habitat (Wiggins et al., 1980). Several species of Trichoptera inhabiting temporary streams have developed special adaptations. This is also the case of Stenophylax species, which are well adapted to replace other limnephilids in temporary waters: all species are univoltine; the eggs are enclosed by a gelatinous matrix that allows them to survive long periods of time out of water; adult or ovarian diapause is present; the burrowing larvae can use the humid substratum during low-water periods, and the pupal cases are buried vertically in the substratum. The Rhyacophilids larvae are probably the caddisflies that are most restricted to conditions of high current-speed (Edington & Hildrew, 1995), occupying, generally, headwaters. Hydropsychids larvae are rheophilic inhabitants of stream rilles (Edington & Hildrew, 1973; Fuller & Mackay, 1980; Osborn & Herricks, 1987; Bonada, 2003). Because of its high dispersion capacity, the survival of both Rhyacophilids and Hydropsychids species during the dry period in Mediterranean streams may involve a high capacity for the reconstruction of colonies from nearby permanent sources (Resh, 1982; 1992).

Though the longitudinal distribution of the Hydropsychids has been largely studied, we think that studies on the longitudinal zonation of endemic or restricted distribution range species, like some present in this catchment, are necessary. The aim of this study was to search for the response of Hydropsychid and Rhyacophilid species present in the river, to the spatial-temporal gradients, as well as to establish their relationship with the water permanence.

METHODS

Study area and sampling methods

The Hozgarganta River basin covers a surface of 245 km² and it extends in a NW - SE direction (Fig. 1). The system drains the Eastern slopes of the mountain range of the Aljibe and after 55 km the watercourse joins the River Guadiaro, near its mouth (Blanco et al., 1991).

The study area included all of the river basins within the limit of the Los Alcornocales Natural Park. The river is born at 160 m a.s.l., at the confluence of the Pasada Blanca canyon and the La Sauceda gorge. The first one flows
through calcareous lands and the second one flows through the Aljibe unit (dominated by sandstone with marl and clay).

The study area is covered by a well preserved Mediterranean forest, in which the cork oak (*Quercus suber*) dominates. Other common species are the Andalusian gall oak (*Quercus canariensis*) and the wild olive (*Olea europaea*). In the humid and shaded canyons there are remains of a subtropical forest, a relict from the tertiary age (*e.g.* *Laurus nobilis*, *Rhododendrum ponticum*) and an arboreal stratum of alder-trees (*Alnus glutinosa*), willows (*Salix atrocinerea*, *Salix pedicellata*), and ash-trees (*Fraxinus angustifolia*) (Jurado-Doña, 2002). In this basin, the land is mostly used for cork oak forest exploitation and extensive ranching.

The flows in successive years show the typical irregularity of the Mediterranean rivers, with dry and wet alternating periods (Fig. 2a). The average annual flow in the period 1980/99 was 1050 m$^3$ s$^{-1}$. In figure 2b we can see that the year when the study was performed (1997) was a wet period.

Another important hydrological feature is the seasonality (Table 1). The maximum registered annual flows are in autumn and early winter, whereas in summer the superficial current disappears, and the water is limited to isolated pools. The average annual period in which there is no superficial flow is 2.65 months (range: 0-6 months, $n = 20$ years).

Nineteen sampling sites were chosen, four in the river’s main channel and the rest on the tributaries (Fig. 1). From November 1996 through December 1997 samplings were carried out in November (1996), February, March, April, June, August, October, and December of 1997. In June, Eca and Ere sampling sites (Table 1) were dry and there-
Table 1. Mean values of the physical-chemical parameters measured in the Hozgarganta basin. Cond: conductivity; Cl\(^{-}\): chloride; Alk: alkalinity; O\(_2\): dissolved oxygen; Temp: temperature; Altit: altitude; Season: seasonality *: 1. permanent flow; 2. ephemeral; 3. intermittent. Inv: invaluable.

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<th>Cl(^{-}) mg/l</th>
<th>Alk meq/l</th>
<th>O(_2) mg/l</th>
<th>Temp °C</th>
<th>Altit a. s. l.</th>
<th>Order</th>
<th>Season</th>
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<th>PO(_4) μgat/l</th>
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fore could not be sampled. In August there was only superficial water in E1, Ecñ, Epbm, Ephh, Eo, Ero, Esa, and Evi sampling sites (Table 1). In December the sampling sites Epbm, Ephh, Ega, and Evi were inaccessible after autumn rains.

A kick net of 0.3 × 0.3 m opening and 0.5 mm mesh size and the same unit of effort in all the localities (three replicates per site) was used for the extraction of the macroinvertebrates. All habitats were sampled along a 50 m riverbed stretch. The samples were fixed using 70% alcohol and later identified to species level in the laboratory. Five Hydropsychidae (Diplectrona felix, Hydropsyche infernalis, H. Siltalai, H. Lobata, and H. iberomaroccana) and two Rhyacophilidae (Rhyacophila munda and Rhyacophila fonticola) species were identified. Because of the taxonomic difficulties in identifying H. Iberomaroccana larvae, this study was complemented with adult records.

For the physical-chemical analysis, samples in February (winter), June (spring), August (summer), and December (autumn) were taken. The average values of the analysed parameters are shown in Table 1.

Figure 2. (X 1000)

a) Registered average annual flows in twenty years in the water gauge station of Jimena de la Frontera. Discharge values × 1000. Caudal medio mensual en el estación de aforo de Jimena de la Frontera. Caudal × 1000. b) Comparison of the monthly average discharges (Qmean) of the temporal series 1981-99 with the registered ones in 1997 (Q97), measures in the water gauge station of Jimena de la Frontera. Comparación del caudal medio mensual (Qmean) de la serie temporal 1981-99 con los registrados en 1997 (Q97), medidas en la estación de aforo de Jimena de la Frontera. Elaborado a partir de la base de datos de la Confederación Hidrográfica del Sur de España.
Data analysis

For the physical-chemical characterisation of the basin a Principal Component Analysis (PCA) on physical-chemical parameters × localities data matrix was performed. To establish the environmental features of the river system, the four most explanatory variables in a previous work in the area (Ruiz et al., 2006) were selected: altitude, conductivity, dissolved oxygen, and temperature.

The ordination method used was selected based on the length of the gradient calculated by Detrended Correspondence Analysis (DCA) (Bonada et al., 2005). Since the first DCA axis has a gradient length of 5.2 standard deviation units, the use of a unimodal ordination technique was justified. To study the relationship between environmental variables and species, a Canonical Correspondence Analysis (CCA) with seasonal abundance data was performed. The significance of CCA axes was tested with the Monte Carlo permutation test (999 unrestricted permutations, $P < 0.005$). All variables used were log-transformed to achieve normality. All the analyses were performed with the PCORD program. To test for the influence of water permanence (seasonality) on the species’ distribution a Kruskal-Wallis test was used.

Figure 3. Classification by Principal Component Analysis based on four selected environmental variables (altitude, conductivity, dissolved oxygen and temperature) of sampling sites of the Hozgarganta catchment. Clasificación de las localidades muestreadas en la Cuenca del río Hozgarganta mediante un Análisis de Componentes Principales basado en cuatro variables (altitud, conductividad, oxígeno disuelto y temperatura).
the thermal preferences of species an adjusted average temperature (AAT) was calculated (Gal-lardo-Mayenco et al., 1998).

RESULTS

The first PCA axis accounted for 52.9% of the variance and shows sampling sites ordered according to altitude and water conductivity, representing a geomorphologic and water permanence gradient, where headstream sites (Em, Eml and Emc) have permanent conditions (Fig. 3). The negative end of this axis was associated to spring samples of downstream sites (E2, Ecñ, E1 and Esa), where high values of both conductivity and temperature were registered. The second PCA axis explained 28.2% of the variance (the cumulative variance was 81.1%). This axis distributed the sampling sites according to the concentration of dissolved oxygen (Fig. 3). We can see that the summer samples from E1 and Epbh are separated from others due to the low value of the dissolved oxygen. Seasonality was of little importance in explaining the sites’ distribution.

The CCA performed from seasonal data showed that the total variance (“inertia”) was 2.75; in the procedure three canonical axes were ob-

Figure 4. Projection of the two first axis of the CCA with seasonal data. Symbol code categories of season (winter (w); spring (sp); summer (s) and autumn (a)) are showed at the end of the acronyms of the sites. Symbol code categories of the variable water permanence (Perm) as follow: 1. permanent; 2. ephemeral; 3. intermittent. + symbol show species. Species codes are explained in Table 2. Sites codes are explained in Table 1.

Proyecci´on de los dos primeros ejes de un CCA con datos estacionales. El c ´odigo de las estaciones del a ˜no (invierno (w); primavera (sp); verano (s) y oto˜no (a)) aparece al final de los acr ´onimos de las localidades de muestreo. El c ´odigo de las categor ´ıas de la variable permanencia del agua en el cauce (Perm) es el siguiente: 1. permanente; 2. ef´ımera; 3. intermitente. El s´ımbolo + representa a las especies. El c ´odigo de las especie es el mismo que aparece en la Tabla 2. El c´odigo de las localidades de muestreo aparece en la Tabla 1.
Table 2. Seasonality influence of the channels (permanent/intermittent/ephemeral) in the species distribution calculated with the Kruskal-Wallis test. Significant \( p \)-values \( (P < 0.005) \) are show in bold. *Influencia de la estacionalidad de los cursos de agua (permanentes/intermitentes/efímeros) en la distribución de las especies, calculada con el test de Kruskal-Wallis. Los valores significativos del ajuste \( (P < 0.005) \) aparecen en negrita.*

<table>
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<td>H. infernalis</td>
<td>Hin</td>
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<td>H. siltalai</td>
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tained. The Monte Carlo test indicated that only the first two axes were significant \( (p = 0.005) \). The first axis explained 28.8 % of the variance; the second one 6.7 %. The cumulative variance was 35.4 %. We obtained that the variables strongly related to the first axis were altitude \( (r = -0.91) \) and conductivity \( (r = 0.88) \), and the temperature was weakly related with the second axis \( (r = 0.18) \). The percentage of unexplained variance was 64.6 %.

Figure 4 shows sites and species distributed in the plane formed by axes 1 and 2. Axis 1 placed the species from left to right according to altitude. *D. felix* and *H. infernalis* occupied headstreams; *R. fonticola* and *H. siltalai* inhabited middle streams and the others were downstream species.

The second factor that determined species’ distribution was water temperature and conductivity. The low half of axis 2 (Fig. 4) grouped most of the high conductivity sites, whereas the top half of this axis discriminated sampling sites with high temperature. Table 2 and figure 5 show the species significantly related \( (p < 0.05) \) to flow permanence. *D. felix* and *H. infernalis* were significantly present in permanent conditions, *H. iberomaroccana* inhabited intermittent sites, while *R. munda*, *R. fonticola*, *H. siltalai*, and *H. lobata* did not show a preference to a particular habitat. *R. fonticola* was mainly associated with the Ero site (intermittent conditions), though it also occupied headwaters (permanent conditions); *H. siltalai* dominated the winter sample from the Emo site (ephemeral conditions). On the other hand, *H. lobata* occupied low reaches, in both intermittent and ephemeral sites (Evi, Ega and E2 stations).

*D. felix* and *H. infernalis* were present in cool water conditions \( (12.1 \text{ and } 12.8^\circ \text{C of AAT, respectively}) \); *H. siltalai* and *H. iberomaroccana-*
were found in sites of moderate temperature (17.7° C and 17.38° C of AAT, respectively); and, *H. lobata* preferred warm conditions (19.4° C of AAT). On the other hand, the Rhyacophilids species showed a similar pattern of thermal preferences, where *R. fonticola* occupied the cool water sites (12.45° C of AAT), while *R. munda* was abundant in warm waters (18.01° C of AAT).

**DISCUSSION**

Numerous studies have pointed out the replacement of species of Hydropsychidae along the watercourse (Camargo, 1992; García de Jalón, 1986; Verneaux & Fassel, 1976; Gallardo-Mayenco *et al*., 1998). This succession may be a consequence of different feeding habits (Voelz & Ward, 1992), metabolic needs (Roux *et al*., 1992), and both differential competitive ability and different mesh size of the species (Tachet *et al*., 1992).

In Mediterranean streams a strong environmental gradient between headwater and lower altitude streams has been observed (Gallardo-Mayenco *et al*., 1998). In our study, the geomorphological gradient is the most important factor explaining the distribution pattern of Hydropsychid species. In the high Hozgarganta basin we have obtained an upstream assemblage, composed of *D. felix* and *H. infernalis*; *H. lobata* inhabits lower reaches in the main river, while *H. iberomaroccana* and *H. siltalai* are widespread throughout the study area, though *H. iberomaroccana* does not reach upstream reaches (e.g. Canutos del Moral, Molino de las Cuevas y del Moro).

Rhyacophila larvae are probably the caddisfly that are most restricted to conditions of high current speed. Their commitment to fast flow rates reflects both the distribution of their food supply and their physiological limitations (Edington & Hildrew, 1995). Their distribution is restricted, almost exclusively, to the headwaters (Tachet *et al*., 2002). In our case, both *Rhyacophila* species display a differentiated distribution pattern. *R. munda* is a wide spectrum ecological species, inhabiting from cool upstream habitats to warm stream ones and the great lower rivers (García de Jalón & González del Tánago, 1986). In the Hozgarganta river, *R. munda* inhabits the principal channel and it is displaced at the headwaters by *R. fonticola*, a species characteristic of springs (Giudicelli & Dakki, 1984) and upstreams (Ruiz *et al*., 2001). A similar result was obtained by García de Jalón & González del Tánago (1986) in the Guadalteba river, where *R. pascoei* displaces *R. munda* at the headwaters. In the same way, Fernández-Aláez *et al*. (2002) found *R. terpsichore* in the upper reaches of the Boeza river, which disappeared in the lower reaches, where ubiquitous species like *R. meridionalis* and *R. relicta* appeared.

Another characteristic of the Mediterranean rivers is the irregularity of their flow, maximum in autumn and spring and minimum in summer (Giudicelli *et al*., 1985) and its high inter-annual variability (McElravy *et al*., 1989), that may imply an inter-annual variability in the conditions of temporality of one site (Del Rosario & Resh, 2000). This determines that the temporary rivers are widely distributed by the climates of Mediterranean type (Gasith & Resh, 1999).

The duration of the dry period has been recognised as an important factor explaining the biological diversity in these rivers (Williams & Hynes, 1976, Abell, 1984, Williams, 1996, Bonada, 2003, Ruegg & Robinson, 2004). Overall, the temporary rivers have fewer species than the permanent ones (Del Rosario & Resh, 2000, Bonada, 2003, Arab *et al*., 2004, Ruegg & Robinson, 2004). In this sense, Bonada (2003) suggests that the low taxonomical richness in ephemeral conditions, and the high difference in richness from permanent sites, would suggest a slow recovery from the last dry period.

Hydropsychids typically build their nets in rapidly-flowing waters (Edington & Hildrew, 1995; Tachet *et al*., 2002) and in mediterranean rivers, they are indicators of riffles conditions, along with Rhyacophilids, (Bonada, 2003). Our study agrees with these results, because *D. felix* and *H. infernalis* are restricted to the upstream reaches; *H. siltalai*. *R. munda* and *R. fonticola* either inhabit intermittent streams or are indifferent to water permanence. In this case, their life cycle is adapted to these environmental conditions, finishing their larval growth in the
wet period (winter-spring). However, *H. iberomaroccanana* and *H. lobata* displayed populations throughout the year, despite the fact that they prefer temporary conditions. Our results are in agreement with those by Gallardo-Mayenco et al. (1998) that, in a study of Hydropsychid species in the Guadaira and Guadalete river basins, point out the opportunistic behaviour of *H. iberomaroccanana* (as *H. punica*), specially due to its capacity to colonise and thrive in temporary habitats. In the Hozgarganta river these species spend the dry season (summer) living in isolated pools, where the absence of superficial flow is noticeable. Numerous studies highlighted the effects of water velocity on the distribution of Hydropsychidae species. Overall, downstream species seem to be more tolerant to low velocity conditions than upstream species (Tachet *et al*., 1992). Hildrew and Edington (1979) showed that *H. pellucidula*, a sister species to *H. iberomaroccanana*, was indifferent or more abundant in low velocity sites. On the other hand, the interruption of the superficial flow does not necessarily imply that of an underground flow between adjacent pools that might remain connected through the aquifer. We suggest that *H. iberomaroccanana* might live in a wide range of current speeds and this allows it to survive in underground-fed pools.

The coexistence of several Hydropsychid species is facilitated by microhabitat partition (Czachorowski, 1989; Harding, 1997), where substrate size and type is important in modulating species’ micro distribution. However, multiple factors operate synergistically over several spatial scales and thus, influence the distribution of the Hydropsychid species (Fairchild & Holomuzki, 2002) in addition to temporal segregation in their life histories (Recasens & Puig, 1987). Hydrological disturbances may also facilitate the coexistence of Hydropsychid species (Resh *et al*., 1990). In our case, *D. felix* and *H. infernalis* were two coexistent species in permanent headstreams. Our data suggests that coexistence is possible, at least partially, by temporal segregation, because *D. felix* was dominant in the spring-summer season while *H. infernalis* was very scarce in summer.

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