

1 Macroinvertebrate community composition and diversity in ephemeral and perennial ponds on
2 unregulated floodplain meadows in the UK

3 M. J. Hill¹, R. G. Death², K. L. Mathers³, D. B. Ryves³, J. C. White³ and P. J. Wood³

4 ¹ Institute of Science and the Environment, University of Worcester, Henwick Grove, Worcester WR2
5 6AJ, UK.

6 ² Institute of Agriculture and Environment, Massey University, Private Bag 11-222, Palmerston North,
7 New Zealand.

8 ³ Centre for Hydrological and Ecosystem Science, Department of Geography, Loughborough
9 University, Loughborough, Leicestershire, LE11 3TU, UK

10

11 **Author for correspondence**

12 Matthew J Hill

13 Institute of Science and the Environment

14 University of Worcester

15 Worcester

16 Worcestershire

17 WR2 6AJ

18 UK.

19

20 Tel: 00 44 (0)1905 542707

21 Email: Matthew.Hill@worc.ac.uk

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23 **Abstract**

24 Ponds are common and abundant landscape features in temperate environments, particularly on
25 floodplains where lateral connectivity with riverine systems persists. Despite their widespread
26 occurrence and importance to regional diversity, research on the ecology and hydrology of temperate
27 ephemeral and perennial floodplain ponds lags behind that of other shallow waterbodies. This study
28 examines the aquatic macroinvertebrate diversity of 34 ponds (20 perennial and 14 ephemeral) on two
29 unregulated riverine floodplain meadows in Leicestershire, UK. Perennial ponds supported nearly
30 twice the diversity of ephemeral ponds. Despite frequent inundation of floodwater and connectivity
31 with other floodplain waterbodies, ephemeral ponds supported distinct invertebrate communities
32 when compared to perennial ponds. When the relative importance of physical and chemical,
33 biological and spatial characteristics was examined, physical and chemical characteristics were found
34 to account for more variation in community composition than biological or spatial variables. The
35 results suggest that niche characteristics rather than neutral colonisation processes dominate the
36 structure of invertebrate communities of floodplain ponds. The maintenance of pond networks with
37 varying hydroperiod lengths and environmental characteristics should be encouraged as part of
38 conservation management strategies to provide heterogeneous environmental conditions to support
39 and enhance aquatic biodiversity at a landscape scale.

40 **Key Words:** community composition, community heterogeneity, connectivity, dry phase duration,
41 hydroperiod, invertebrate, species richness

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48 **Introduction**

49 Floodplain landscapes are sites of exceptionally high aquatic, semi-aquatic and terrestrial diversity
50 (Ward et al. 1999; Helfield et al. 2012) which may be strongly influenced by lateral connectivity to
51 lotic ecosystems (Tockner et al. 2000; Starr et al. 2014). The flooding of riverine landscapes creates
52 and maintains a variety of aquatic habitats and typically results in a network of hydrologically
53 connected perennial and ephemeral waterbodies at a range of successional stages (Paillex et al. 2013).
54 However, due to anthropogenic flow regulation, embankment and channelization to reduce flood risk
55 and to protect infrastructure and agricultural activities on the floodplain, many rivers are
56 hydrologically disconnected from their floodplain along most of their course (Nilsson et al. 2005;
57 Paillex et al. 2013). This has resulted in a long term trend of terrestrialization of floodplain habitats
58 compounded by agricultural expansion and urbanisation leading to a reduction in freshwater
59 biodiversity and habitat (Tockner & Stanford 2002; Reckendorfer et al. 2006).

60 Ponds located on traditionally managed floodplains can provide important habitats for a wide range of
61 unique flora and fauna (Shiel, et al. 1998; Williams et al. 2008). Floodplain ponds support diverse
62 aquatic habitats and often represent locations of high alpha (site), beta (between ponds) and gamma
63 (regional) diversity (Gergel 2002). They are common and abundant aquatic habitats globally
64 (Williams 1997; Boix et al. 2016) often occurring in pond networks but they have been poorly studied
65 in most regions compared to other freshwater habitats (Gergel 2002; Williams 2006). Many
66 floodplain ponds are ephemeral (they experience recurrent drying; Williams et al. 2001), and are often
67 characterised by a gradient of permanence (hydroperiod), from those containing water for a few
68 months through to those with perennial surface water. Floodplain ponds therefore have the potential to
69 be strongly controlled by colonisation dynamics, but may equally be driven by local habitat conditions,
70 particularly if some ponds dry while others remain wet.

71 The physical and chemical conditions of ephemeral ponds are demanding for biota and often become
72 extreme as the pond dries and aquatic habitat is lost (Williams 1996; Williams 2006; Bagella et al.
73 2010). Due to the potentially wide range of conditions they experience, ephemeral ponds have been
74 shown to be important habitats for a diverse range of macroinvertebrate taxa adapted to and able to

75 exploit the conditions they offer (Bazzanti et al. 2010). Although ephemeral ponds often support a
76 lower taxonomic richness than perennial ponds, they may support a high richness of ‘rare’ and
77 endemic taxa (Nicolet et al. 2004; Armitage et al. 2012) and in some cases support a greater number
78 and proportion of rare taxa than perennial ponds in close geographical proximity (Collinson et al.
79 1995; Della Bella et al. 2005). Fish typically occur in low abundances or are absent from ephemeral
80 ponds as they cannot withstand desiccation which may greatly reduce predation pressure on
81 invertebrates (although high predation pressure may still occur from other vertebrates and
82 invertebrates e.g., Amphibia, Coleoptera and Crustacea; Brendonck et al. 2002). The absence of fish
83 may also increase the abundance/richness of open water taxa and other fauna that may be
84 outcompeted in perennial ponds (Bronmark & Hansson 2005; De Meester et al. 2005).

85 There has been a recent drive to re-connect rivers with their floodplains and to rehabilitate and restore
86 aquatic habitats on the floodplain to support faunal and floral diversity (Buijse et al. 2002;
87 Reckendorfer et al. 2006; Paillex et al. 2015). However, debate surrounds the relative importance of
88 local habitat (referring to the physical, chemical and biological characteristics of individual ponds)
89 and regional (connectivity/isolation: the spatial configuration of ponds) variables in determining pond
90 community composition (Vanschoenwinkel et al. 2007). Although the physical and chemical
91 characteristics of ponds have been considered in some detail (Hinden et al. 2005; Hassall et al. 2011),
92 most have largely ignored the relative role of regional variables in influencing community
93 composition (Van de Meutter et al. 2007; Heino et al. 2014). Metacommunity theory provides a
94 theoretical framework to partition the mechanisms that may underlie biological distributions in a pond
95 network (Leibold et al. 2004; Vanschoenwinkel et al. 2007). A metacommunity is defined as ‘a set of
96 local communities that are linked by dispersal of multiple potentially interacting species’ (Leibold et
97 al. 2004: 602) where communities are located on a continuum from those dominated entirely by
98 regional colonisation dynamics, to those where niche differentiation based on local habitat conditions
99 dominate. Four general community types can therefore be recognised; 1) *patch dynamics* - numerous
100 homogenous patches are present in which the driving force of community structure is a trade-off
101 between competitive ability and dispersal (Leibold et al. 2004); 2) *species sorting* - species distribute

102 amongst heterogeneous patches based on their ability to specialize within particular abiotic niches
103 (Cottenie et al. 2003; Vanschoenwinkel et al. 2007); 3) *mass effects* - dispersal drives community
104 composition. Different patches experience different conditions at a given time and dispersal of
105 individuals between patches is frequent, creating source-sink relationships. Local extinctions of
106 individual species can be prevented by dispersal from patches where they are good competitors
107 (Heino et al. 2014); and 4) *the neutral view* - which assumes species are functionally equivalent and
108 distribute amongst patches at random (Leibold et al. 2004).

109 To investigate the potential local and regional drivers of pond community composition and diversity
110 we quantified the macroinvertebrate diversity and community structure of ephemeral and perennial
111 ponds located in largely unregulated floodplain meadows. We examined whether spatial proximity
112 (neutral processes) or local environmental variables (niche processes) dominated macroinvertebrate
113 community composition among the ephemeral and perennial ponds.

114

115 **Methods**

116 Study area and sites

117 Ponds are defined here as small lentic water bodies between 25 m² and 2 ha in area, frequently less
118 than 2 m deep, which normally hold water for at least 4 months of the year (Williams et al. 2010). A
119 comprehensive examination of 34 ponds was undertaken on two largely unregulated floodplain
120 meadows adjacent to the River Soar, Leicestershire, UK: Cossington Meadow (25 ponds, ~86 ha, lat:
121 52.715621 long: -1.116947) and Loughborough Big Meadow (9 ponds, ~60 ha, lat: 52.789178 long: -
122 1.116947). Both meadows are located in nature conservation areas and are naturally inundated by
123 water from the River Soar during the winter and early spring each year. Fluvial gravel and sand were
124 historically quarried from Cossington Meadow, but since 2004 it has been a protected nature reserve
125 supporting a variety of floodplain meadow, woodland and freshwater habitats (perennial and
126 ephemeral ponds, lakes and ditches), all in close proximity to the River Soar. The majority of the
127 larger ponds and lakes are of anthropogenic origin (relicts of quarrying) but since their creation,

128 limited direct management has been undertaken and they are minimally affected by low density
129 pastoral agriculture associated with traditional floodplain meadow systems. Loughborough Big
130 Meadow is part of a Site of Special Scientific Interest and is one of the few remaining traditional
131 floodplain Lammas meadows in the UK. Lammas refers to a particular type of land tenure. During the
132 crop-raising period (February to August) the land owners divide the meadow into sections and sell the
133 rights to the hay crop to local farmers. Once the hay crop has been gathered the land becomes subject
134 to the rights of common grazing (mid-August - February). The study took place during 2012 and was
135 characterised by drought conditions at the start followed by a period of sustained high rainfall (Marsh
136 et al., 2013). In some regions of the UK this resulted in significant variability in water levels and
137 wetting and drying of temporary ponds (Jeffries 2015). However, the lowland location of the ponds in
138 this study meant that at the start of the sampling programme the majority of pond basins were wet and
139 although water levels were highly variable, the total number of inundation events (floods) and
140 duration that the basins were dry (hydroperiod) was comparable to average conditions.

141

142 Aquatic macroinvertebrate sampling

143 The ponds studied comprised two groups: (i) 20 perennial ponds - water bodies which contained water
144 all year round and; (ii) 14 ephemeral ponds - ponds which became dry (dry phase varied from 3-6
145 months) at least once during the study period (Jan 2012 - Dec 2012). Floodwater recharge from the
146 River Soar was the primary driver of hydroperiodicity for the ephemeral ponds studied. Aquatic
147 macroinvertebrate samples were collected on three occasions from each pond corresponding to spring,
148 summer and autumn seasons. The total number of samples taken was 87 (perennial n=60, ephemeral
149 n=27). All temporary ponds dried at least once during the sampling period and were not sampled
150 during the dry phase. In this study the sampling strategy of fixed timed macroinvertebrate collections
151 was deemed not suitable to examine diversity within the small and ephemeral ponds where the wetted
152 area varies seasonally (Armitage et al. 2012). The strategy was therefore modified to obtain
153 representative samples from all sites whilst ensuring that the small freshwater habitats/communities
154 were not adversely affected by the sampling (Armitage et al. 2012). The sampling time allocated to

155 each pond was proportional to its surface area up to a maximum of 3 minutes (Biggs et al. 1998). The
156 maximum sampling time of 3 minutes was used for ponds with a surface area $>50 \text{ m}^2$; for smaller
157 ponds 30 seconds of sampling for every 10 m^2 surface area was employed. A standard pond net (mesh
158 size, 1 mm) was used to sample aquatic macroinvertebrate taxa. The total sampling time designated to
159 each individual pond was divided equally between the mesohabitats present (open water, emergent
160 macrophytes and submerged macrophytes). If one mesohabitat dominated the pond, sampling time
161 was further divided to reflect this; for example, in a pond with 3 mesohabitats sampling time was
162 divided by 4 – one from each mesohabitat with an additional sample from the dominant mesohabitat
163 (Biggs et al. 1998). In addition, an inspection of hard surfaces or larger substrate (e.g., rocks and large
164 floating leaves) for aquatic macroinvertebrates was undertaken for 1 minute at each site. In the
165 laboratory, aquatic macroinvertebrate samples from each habitat were processed and preserved in 70%
166 industrial methylated spirits prior to identification. Taxa were identified to species level except,
167 Diptera larvae, Planariidae, and Hydrachnidiae which were identified to order or family level and
168 Oligochaeta and Collembola were recorded as such. The macroinvertebrate taxa with UK
169 conservation designations were identified using the extensive list provided by the JNCC (JNCC 2015).

170

171 Environmental data collection

172 The following local environmental parameters were measured at each site prior to macroinvertebrate
173 sampling: surface area (wetted area: m^2), mean water depth (cm), the percentage of the pond margin
174 and pond surface shaded by overhanging vegetation, the presence of fish (0/1 as a dummy variable);
175 and dry phase length (duration in months between Jan-Dec 2012 that the pond was dry). Conductivity
176 ($\mu\text{S cm}^{-1}$), pH (Hanna Instruments - HI198311 and HI98127) and dissolved oxygen (DO mg l^{-1})
177 (Mettler Toledo DO Meter SG6) were measured at the margin of each site using portable meters. The
178 occurrence and proportion (% of surface area) of mesohabitats within each pond was recorded.
179 Regional environmental variables; Pond connectivity - number of waterbodies hydrologically
180 connected to a sample site (e.g., through rivulets or overland flooding) and pond proximity - the
181 number of other fresh waterbodies within 500 m (Vanschoenwinkel et al. 2007; Waterkeyn et al.

182 2008), were recorded through visual inspection (walking extensively around each site during each
183 season to identify nearby perennial and ephemeral ponds and through the use of aerial imagery
184 provided by Google Earth Software (Goole Earth 2015). Every attempt was made to record all
185 waterbodies within 500 m of each meadow pond site, however, small temporary ponds can be difficult
186 to identify through visual inspection and aerial images and it is therefore acknowledged that a small
187 number of temporary ponds may have been overlooked.

188

189 Statistical analyses

190 Aquatic macroinvertebrate diversity was examined across the floodplain meadow landscape (gamma
191 diversity) and for individual ephemeral and perennial ponds (alpha diversity). Macroinvertebrate
192 abundance and taxon richness were calculated for each mesohabitat and pond site (mesohabitat and
193 seasonal data for each pond site were combined to provide a total measure of diversity for each study
194 site) using PRIMER 6 (Clarke and Gorley 2006). Ecological diversity is heavily affected by the
195 sample size and sampling procedures (McCabe & Gotelli 2000). As a result, rarefaction (Hulbert 1971)
196 was undertaken in PRIMER 6 to estimate species richness for each mesohabitat and pond site for a
197 given number of individuals drawn randomly from a sample (McCabe & Gotelli 2000). The least
198 abundant sample had 28 individuals; as a result, 28 individuals were randomly sampled from each
199 mesohabitat and pond site and the rarefied species richness was recorded.

200 The statistical significance of differences in faunal diversity among the ephemeral and perennial pond
201 types and mesohabitats (open water, emergent and submerged macrophytes) were examined using a
202 nested analysis of variance (nested ANOVA) with Bonferonni *post hoc* tests. Pond type and
203 mesohabitat were included as fixed effects and site was nested within pond type as a random effect.
204 Differences in the dispersal characteristics and functional feeding groups of macroinvertebrate
205 communities between ephemeral and perennial ponds were examined using a non-parametric
206 ANOVA (Kruskal-Wallis test). Dispersal and functional feeding traits assigned to individual
207 macroinvertebrate taxa follow the classification of Tachet et al. (2003) and Merritt and Cummins
208 (1996). Variability in physical and chemical parameters between pond sites were examined using one-

209 way analysis of variance (ANOVA). The data were examined to ensure they complied with the
210 underlying assumptions of parametric statistical tests (e.g., normal distributions) and abundance data
211 were \log_{10} transformed where required. All univariate analyses were undertaken in IBM SPSS
212 Statistics (version 21, IBM Corporation, New York). The Waikato Environment for Knowledge
213 Analysis (WEKA) machine learning software (version 3.6.1) was used to construct regression trees to
214 predict taxa richness of the ponds from the collected environmental data (Witten et al. 2011). A
215 regression tree was generated with the M5P option and 10% cross validation in WEKA (Quinlan 1993;
216 Witten & Frank 2000).

217 The variability of macroinvertebrate communities was described using MVDISP in PRIMER 6
218 (Clarke & Gorley 2006) to compare the multivariate dispersion (compositional variability) of
219 communities in ephemeral and perennial ponds. Community heterogeneity between ephemeral and
220 perennial pond sites was statistically examined using Analysis of Similarity (ANOSIM) in PRIMER 6
221 (Clarke & Gorley 2006). Prior to ANOSIM analysis, faunal-abundance data were $\log(X+1)$
222 transformed. The PRIMER 6 program RELATE (a mantel-type test) was used to examine the
223 relationship between the aquatic macroinvertebrate community dissimilarity and spatial distance
224 (meters) and environmental distance (Euclidean). RELATE tests the significance of a Spearman's
225 rank correlation between two distance matrices (Bray-Curtis community dissimilarity and geographic
226 distance between study pond sites). To test the association between macroinvertebrate taxa and pond
227 type and identify indicator taxa of ephemeral and perennial ponds Indicator Value analysis (IndVal)
228 (Dufrière & Legendre 1997) was undertaken in R (R Development Core Team 2013).

229 The associations between macroinvertebrate community composition and environmental variables
230 (local and regional) were assessed using Redundancy Analysis (RDA) implemented in the programme
231 CANOCO (Version 4.5; ter Braak & Šmilauer 2002). Due to natural variability in macroinvertebrate
232 community assemblages, seasonal faunal data from individual pond sites were combined and mean
233 values of environmental variables calculated. Prior to analysis, environmental parameters were \log_{10}
234 transformed (except for pH) to reduce the influence of skew in the data set and overcome the effect of
235 their physical units (Legendre & Birks 2012). Faunal-abundance data were Hellinger transformed

236 prior to analysis (Legendre & Gallagher 2001). A forward selection procedure, using a random
237 Monte-Carlo permutations test (999 random permutations) with Bonferroni correction was employed
238 to determine the significance of the relationship between the environmental variables and
239 macroinvertebrate composition. Only physical and chemical parameters significantly influencing the
240 faunal data ($p < 0.05$ before Bonferroni correction) were included in the final model.

241 Variance partitioning analysis was used undertaken using CANOCO 4.5 to examine the amount of
242 variation in macroinvertebrate community assemblage that can be explained by local (physical and
243 chemical or biological) and regional (spatial) variables (Borcard et al. 1992). Only environmental
244 parameters from the RDA identified to significantly influence macroinvertebrate community
245 composition were used in the variance partitioning analysis. The total percentage of variance
246 explained by the RDA was partitioned into unique contribution (percentage of variance explained by
247 each individual group of environmental variables), and common contributions (variation explained by
248 a combination of groups of environmental variables) using partial RDAs (Borcard et al. 1992;
249 Vanschoenwinkel et al. 2007).

250

251 **Results**

252 Environmental characteristics

253 Environmental conditions recorded among ephemeral and perennial ponds from the two meadow sites
254 were highly variable (Table 1). Perennial ponds were on average twice as deep (ANOVA $F_{1,33} = 37.65$,
255 $p < 0.001$), had higher pH (ANOVA $F_{1,33} = 11.12$, $p < 0.002$) and conductivity (ANOVA $F_{1,33} = 18.28$,
256 $p < 0.001$) than ephemeral ponds. The proportion (%) of the pond covered by emergent macrophytes
257 was nearly four times greater for ephemeral ponds compared to perennial ponds (ANOVA $F_{1,33} = 5.52$,
258 $p < 0.025$) (Table 1). Surface area, surface water shaded, pond margin shaded, submerged macrophyte
259 cover and dissolved oxygen did not differ significantly between ephemeral and perennial ponds
260 ($p > 0.05$). Fish were present in 19 perennial ponds but were absent from all ephemeral ponds.

261 Macroinvertebrate diversity

262 Across the two floodplain meadows, a total of 173 taxa were recorded within 16 orders and 56
263 families from the ephemeral (93 taxa) and perennial ponds (164 taxa; see Supplementary Material
264 Appendix 1 and Appendix 2 for full list of taxa). Macroinvertebrate taxon richness varied widely
265 among pond sites ranging from 5 (ephemeral pond) to 73 (perennial pond) taxa. Macroinvertebrate
266 assemblages within ephemeral and perennial ponds were dominated taxonomically by Coleoptera (Fig.
267 1). On average, hemipteran taxa constituted a much higher proportion of the species richness recorded
268 in perennial ponds (>21%) than ephemeral ponds (<10%). In contrast, Diptera and Crustacea taxa
269 formed, on average, a greater proportion of the taxa richness in ephemeral than perennial ponds (Fig.
270 1). The taxa most widely distributed across the meadow pond sites were Chironomidae larvae (32
271 ponds), Oligochaeta (30 ponds) and *Crangonyx pseudogracilis* (28 ponds). A total of 9
272 macroinvertebrate taxa were only recorded in the ephemeral ponds (*Galba trunculata*, *Libellula*
273 *quadrifasciata*, *Limnephilus auricula*, *Limnephilus centralis*, *Limnephilus griseus*, *Gerris gibbifer*,
274 Elmidae larvae, *Helophorous dorsalis* and *Paracymus scutellaris*).

275 Perennial ponds supported nearly three times the mean taxon richness (ANOVA $F_{1, 105} = 21.75$;
276 $p < 0.001$) and twice the rarefied taxon richness (ANOVA $F_{1, 81} = 11.20$; $p < 0.001$) compared to
277 ephemeral ponds (Table 2). Mean macroinvertebrate abundance (ANOVA $F_{1, 129} = 5.49$; $p < 0.05$) in
278 ephemeral ponds was 20% of that in perennial ponds (Table 2). A significant difference in the number
279 of taxa (ANOVA $F_{2, 109} = 9.77$; $p < 0.001$), rarefied taxa richness (ANOVA $F_{2, 109} = 3.08$; $p < 0.05$) and
280 marginally significant difference in abundance (ANOVA $F_{2, 109} = 3.07$; $p < 0.051$) was observed among
281 the meadow ponds when individual mesohabitat units were considered. Macroinvertebrate abundance
282 was typically greater amongst emergent macrophytes (Fig. 2a). Macroinvertebrate richness and
283 rarefied richness were higher within submerged macrophytes and emergent macrophytes than open
284 water for all ponds (Fig. 2b; 2c). The regression tree analysis yielded a single regression equation:
285 $Taxa\ number = 6.312 * Log\ area + 7.6575 * pH - 43.2272 * Log\ Hydroperiod\ dry\ months + 7.1705 *$
286 $Log\ emergent\ macrophytes - 29.4961$. The cross validated correlation coefficient of 0.86, indicating
287 that the regression equation was a good predictor of taxa number.

288 When functional feeding groups were examined, a greater proportion of the macroinvertebrate
289 community were scrapers and deposit feeders in ephemeral ponds, whilst piercers constituted a
290 greater proportion of the communities recorded in perennial ponds (Fig. 3a). There were a greater
291 proportion of non-predatory taxa recorded in ephemeral ponds (mean: 73%) than perennial ponds
292 (mean: 58% Kruskal-Wallis $p < 0.05$). The proportion of passively and actively dispersing taxa did not
293 differ statistically between the two pond types ($p > 0.05$) (Fig. 3b).

294 Macroinvertebrate community composition was significantly different for ephemeral and perennial
295 ponds (ANOSIM $R = 0.581$, $p < 0.005$). Ephemeral meadow ponds had a higher multivariate dispersion
296 (1.56) than perennial ponds (0.73) indicating that ephemeral ponds displayed greater community
297 heterogeneity than those of perennial ponds (Table 2; Fig. 4). Macroinvertebrate taxa identified as
298 indicator species for ephemeral and perennial meadow ponds are presented in Table 3.

299 Macroinvertebrate - environment associations

300 RDA indicated that five environmental variables (connectivity, pond proximity, pond surface area,
301 submerged macrophyte coverage and the dry phase duration) had a significant influence on
302 community composition (Fig. 4; Monte Carlo Tests $F = 3.33$ $p < 0.005$) with all axes explaining 45.8%
303 of the assemblage variance. A clear distinction between ephemeral (towards the bottom right) and
304 perennial ponds (far left and top) was apparent in the RDA biplot (Fig. 4). A cluster of 12 perennial
305 ponds directly connected to each other and the River Soar plotted on the far left of axis 1 (Fig. 4a).
306 These ponds were inundated twice by floodwater from the River Soar during the sampling period. The
307 other perennial meadow ponds typically had larger surface areas (Fig. 4a). The seasonal drying of the
308 pond basin ($F = 3.77$ $p < 0.01$) was identified to be a key parameter structuring macroinvertebrate
309 composition among ephemeral meadow ponds (Fig. 4a). In addition, ephemeral ponds were associated
310 with reduced pond proximity. The highest taxon richness was typically associated with greater surface
311 area ($F = 2.3$ $p < 0.01$), pond connectivity and pond proximity to other waterbodies ($F = 4.12$ $p < 0.01$)
312 whilst the lowest richness was associated with longer dry phases (Fig. 4b).

313 Local and regional environmental factors

314 Variance partitioning indicated a greater influence of local physical and chemical variables on
315 community composition (10.8% of total variance) compared to spatial (4.5%) or biological variables
316 (4.1%; Fig. 5) among the meadow ponds studied. A combination of physical, chemical and spatial
317 variables provided the greatest explanation of community composition (11.8%) among the meadow
318 ponds. Community composition was more different between ponds that were further apart (ρ : 0.507
319 $p < 0.001$) or that differed in local habitat conditions (ρ : 0.586 $p < 0.001$).

320

321 **Discussion**

322 Macroinvertebrate diversity

323 Perennial meadow ponds supported nearly twice the number of macroinvertebrate taxa compared to
324 ephemeral ponds, based on rarefied taxa richness. Several other studies have reported perennial ponds
325 support significantly greater richness than ephemeral ponds in both Temperate and Mediterranean
326 landscapes (Collinson et al. 1995; Nicolet 2001; Della Bella et al. 2005). However, in contrast to the
327 meadow ponds in this study, previous studies have reported more actively dispersing taxa in
328 ephemeral than perennial ponds (Nicolet 2001; Nicolet et al. 2004). The greater proportion of less-
329 mobile taxa in these UK ephemeral ponds may reflect the frequent floodplain inundation, and mixing
330 of water across the floodplain (high connectivity), which would facilitate the migration of passively
331 dispersing taxa from perennial to ephemeral pond habitats (Nicolet et al. 2004). The greater
332 proportion of non-predatory macroinvertebrate fauna recorded from ephemeral ponds most likely
333 reflects the short hydroperiod (typically 6 months). This probably reduced the colonisation potential
334 and occurrence of some larger, longer-lived predators (e.g., Coleoptera, Odonata, fish) which
335 typically have generation times greater than the hydroperiod of the ephemeral ponds (Bilton et al.
336 2001; De Meester et al. 2005; Williams 2006). However, other studies have demonstrated that highly
337 mobile aquatic predators will commonly colonize temporary ponds in spring and disperse to perennial
338 ponds during the summer, with some Coleoptera remaining in damp patches within temporary pond
339 basins after open water has receded and may only disperse more widely when the basin has dried
340 completely (Davy-Bowker et al. 2002).

341 When placed in a national context, the average richness of ephemeral meadow ponds in this study (19
342 taxa) was lower than that recorded in a UK wide study of temporary ponds (25 taxa: Nicolet et al.
343 2004) and elsewhere in the UK (Bilton et al. 2009; Armitage et al. 2012). However, direct comparison
344 is not straightforward as taxonomic resolution, habitat quality and sampling strategies differ between
345 the studies. Macroinvertebrate diversity of ponds in this study is almost certainly significantly higher
346 since Diptera were only resolved to family level. In addition, semi-aquatic and terrestrial riparian
347 fauna (Carabidae and Staphylinidae) that frequently utilise pond basins during the dry phase (Lott
348 2001) were not recorded here or in other studies of ephemeral ponds (Della Bella et al. 2005; Dell et
349 al. 2014) and clearly represents an underestimation of their contribution to biodiversity (Collinson et
350 al. 1995; Drake, 2001).

351 Several gastropod taxa (*L. palustris*, *R. balthica* and Physidae) and the juvenile life stages of
352 Dytiscidae (Coleoptera) and Corixidae were identified as indicator taxa of perennial ponds in this
353 study. The Gastropoda, *L. palustris*, *R. balthica* and Physidae, were widely distributed in perennial
354 ponds, but occurred infrequently in ephemeral ponds as they cannot withstand prolonged desiccation
355 (Nicolet 2001; Della Bella et al. 2005). In contrast, the gastropod *A. leucostoma* was common in
356 ephemeral ponds and can survive desiccation by burrowing into sediments and entering a state of
357 diapause (Bratton 1990). Similarly, the larvae of Dytiscidae and Corixidae were largely confined to
358 perennial ponds since they are unlikely to survive the dry phase within ephemeral pond basins.
359 Although not exclusive to the ephemeral ponds, *Hesperocorixa sahlberghi* was also identified as an
360 indicator of ephemeral ponds. *H. sahlberghi* frequently colonises densely vegetated habitats
361 (emergent macrophyte coverage was greater in ephemeral ponds) and may have also benefited from
362 the absence of predatory fish (Savage 1989).

363 Macroinvertebrate community composition

364 Community composition was strongly associated with habitat characteristics (45.8% of variance was
365 explained); although the strength was lower than for other studies of small pond or rock-pool
366 communities (e.g., Vanschoenwinkel et al. 2007), reflecting the effect of local (e.g. physical and
367 chemical factors) and regional (i.e. connectivity / proximity) parameters in the analysis (Florencio et

368 al. 2014). The community composition recorded in this study was more strongly linked with the
369 physical and chemical characteristics of the pond rather than biological or regional drivers. Local
370 environmental variables also explained more of the variance in macroinvertebrate community
371 composition for ephemeral ponds than how the ponds were distributed in space in South Africa
372 (Vanschoenwinkel et al. 2007) and Donana National Park, Spain (Florencio et al. 2014). Connectivity
373 between ponds can have a homogenizing effect on community structure, increasing diversity as taxa
374 are able to disperse more freely (Cottenie et al. 2003), although other studies have shown this effect to
375 be stronger for passively dispersing taxa than for active dispersers (Vanschoenwinkel et al. 2007). In
376 the current study, more distant ponds did have more dissimilar communities, but spatial factors were
377 of secondary importance to the local habitat (Cottenie et al. 2003; Cottenie & De Meester 2003).

378 If these ponds were placed into the metacommunity framework, the heterogeneity of the habitats and
379 macroinvertebrate communities violate the key assumptions for patch dynamics to apply (assumes
380 that habitat patches are identical; Vanschoenwinkel et al. 2007). A combination of mass effects
381 (connectivity and pond proximity) and species sorting (physical, chemical and biological; Leibold et
382 al. 2004) would probably most effectively explain the macroinvertebrate assemblages (Cottenie et al.
383 2005; Vanschoenwinkel et al. 2007; Ng et al. 2009). Spatial factors (mass effects) promote the
384 dispersal and colonization of invertebrates within the metacommunity but it is the variation in local
385 physical and chemical factors (species sorting) that regulates and controls community composition
386 (Cottenie et al. 2003; Cottenie & De Meester 2003).

387 The greater importance of local variables over regional variables may explain the high community
388 heterogeneity recorded between ephemeral and perennial ponds (Collinson et al. 1995; Della Bella et
389 al. 2005). While high connectivity (floodwater inundation) promotes the dispersal of invertebrates
390 between ephemeral and perennial ponds, it is the local pond conditions (e.g., hydroperiod, wetted area,
391 depth, emergent macrophyte coverage) which sorts and structures the communities. However, the
392 results of this study also indicates many taxa from ephemeral ponds also occur in perennial ponds
393 (Bazzanti et al. 2003; Nicolet et al. 2004; Bilton et al. 2009). Many taxa common to both pond types
394 were generalists, including several Diptera families (Culicidae and Tipulidae spp.) which have the

395 prerequisite traits for successful colonisation and development in ephemeral waterbodies including;
396 rapid development, rapid recolonization via aerial dispersal and the ability of some larvae to persist in
397 damp sediments (Drake 2001). The high density and hydrological connectivity (regular inundation) of
398 ephemeral and perennial ponds on the floodplains would have increased the opportunity for passive
399 dispersal events and allowed many perennial pond taxa to colonise the ephemeral ponds on the
400 floodplain (Nicolet et al. 2004).

401 High connectivity between the river and floodplain can lead to short-term reductions in species
402 richness in systems where large floods disturb the wetland habitats and reset successional trajectories
403 (Bornette et al. 1998; Reckendorfer et al. 2006; Tockner et al. 2010). The floodplain meadows in the
404 current study were not subject to any high magnitude floods during the study period and the high
405 species richness and community heterogeneity among ponds reflects the range of successional stages
406 present, and the gradual re-filling and re-wetting of the lentic (and potentially hyporheic) habitats
407 which facilitate the dispersal of macroinvertebrates and resources (Lake et al. 2006; Starr et al. 2014;
408 Paillex et al. 2015). The absence of erosive floodwaters was also important in structuring the
409 macrophytes within both the perennial and ephemeral ponds. Aquatic macrophytes were found to be
410 important determinants of assemblage and diversity in this and in other studies (Bazzanti et al. 2010;
411 Florencio et al. 2014). This reflects the importance of macrophytes as structurally diverse and
412 complex habitats with abundant niches for aquatic invertebrates, their capacity to serve as refugia
413 from predation, provide sites for oviposition and provide an abundance of trophic resources (Bazzanti
414 et al. 2010).

415 Conservation of floodplain meadow ponds

416 Perennial and ephemeral floodplain meadow ponds provide a valuable and important habitat for
417 aquatic macroinvertebrates, supporting a wide diversity of fauna at an alpha and gamma scale and a
418 number of taxa of conservation interest (Armitage et al. 2012). Despite this, there is limited formal or
419 direct legislative protection (e.g., from the Water Framework Directive or the Habitats Directive,
420 Hassall et al. 2016) of ephemeral ponds in temperate regions at a European scale (Williams et al. 2001;
421 Nicolet et al. 2004). However, it is important to recognise that at a national scale in the UK,

422 ephemeral and perennial ponds may be protected via designation as a priority habitat (BRIG 2008). In
423 addition, the meadow ponds in this study were located in established nature reserves which indirectly
424 provided protection for the ponds and help maintained a high density of ephemeral and perennial pond
425 habitats (and high macroinvertebrate diversity).

426 Natural inundation of the floodplain and riparian meadows would have historically been typical of
427 many temperate zone lowland systems prior to land drainage, agricultural improvement and river
428 regulation. Reconnecting the river with its floodplain will provide significant opportunities to re-
429 naturalize floodplains (Reckendorfer et al. 2006; Castella et al. 2015), however many temperate rivers
430 have poor water quality and polluted floodwater may significantly reduce taxonomic diversity of
431 freshwater bodies on the floodplain (Tockner & Stanford 2002). Strategies to improve river water
432 quality should be implemented alongside river-floodplain reconnection to take advantage of the
433 bioremediation (nutrient storage and processing) potential of floodplain water bodies. However, care
434 is also required to ensure that floodplain wetland and pond restoration is not compromised or
435 prevented due to pre-existing poor river water quality. The reconnection of the channel to the
436 floodplain is will also provide additional refuge habitat for many floral and faunal taxa, potentially
437 increasing ecosystem resilience and the long-term sustainable management of floodplain waterbodies.

438 Results of this study indicate that pond biodiversity conservation on floodplains should primarily
439 focus on improving local habitat quality and diversity. For example, management practices should
440 aim to maintain a diverse array of ephemeral and perennial ponds on floodplains (encompassing the
441 full hydrosere successional sequence) with varying hydroperiod lengths and environmental conditions
442 (Biggs et al. 1994; Williams et al. 2003; Bilton et al. 2009) in order to provide a wide range of niches
443 for invertebrate taxa to utilise. However, wherever possible pond connectivity should be increased on
444 floodplains to provide greater opportunities for macroinvertebrate dispersal and colonisation
445 (Williams et al. 2008). The creation of new ephemeral and perennial pond basins on the floodplain
446 will increase connectivity and dispersal potential between the river and existing floodplain
447 waterbodies (including ponds) and will also provide new high quality sites for macroinvertebrate taxa
448 to utilise. Further, where appropriate the excavation of small rivulets (channels) may increase

449 connectivity between individual ponds and enhance dispersal potential. Quantifying aquatic
450 macroinvertebrate diversity and distribution on unregulated (semi)natural floodplain meadows (across
451 all waterbody types) potentially provides important information regarding the reference conditions for
452 these increasingly rare systems. This is an essential pre-requisite for the ongoing conservation of
453 existing sites and the future restoration and, where both socially acceptable and possible, the re-
454 connection of rivers to their floodplains.

455

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462

463 **References**

- 464 Armitage, P. D., A. Hawczak, and J. H. Blackburn. 2012. Tyre track pools and puddles -
465 Anthropogenic contributors to aquatic diversity. *Limnetica* 42: 254-263
- 466 Bagella, S., S. Gascon, M. C. Caria, J. Sala, M. A. Mariani, and D. Boix. 2010. Identifying key
467 environmental factors related to plant and crustacean assemblage in Mediterranean temporary ponds.
468 *Diversity Conservation* 19: 1749-1768
- 469 Bazzanti, M., V., D. Bella, and M. Seminara. 2003. Factors affecting macroinvertebrate communities
470 in astatic ponds in Central Italy. *Journal of Freshwater Ecology* 18: 537-548
- 471 Bazzanti, M., C. Coccia, and M. Giuseppina Dowgiallo. 2010. Microdistribution of
472 macroinvertebrates in a temporary pond of Central Italy: taxonomic and functional
473 analyses. *Limnologica-Ecology and Management of Inland Waters* 40: 291-299

474 Biggs, J., A. Corfield, D. Walker, M. Whitfield, and P. Williams. 1994. New approaches to pond
475 management. *British Wildlife* 5: 273-287

476 Biggs, J., G. Fox, P. Nicolet, M. Whitfield, and P. Williams. 2001. Dangers and opportunities in
477 managing temporary ponds. *Freshwater Forum* 17: 71-80

478 Biggs, J., G. Fox, M. Whitfield, and P. Williams. 1998. A guide to the methods of the National Pond
479 Survey. Pond Action, Oxford

480 Bilton, D. T., A. Foggo, and D. Rundle. 2001. Size permanence and the proportion of predators in
481 ponds. *Archiv fur Hydrobiologie* 151: 451-458

482 Bilton, D. T., L. C. McAbendroth, P. Nicolet, A. Bedford, S. D. Rundle, A. Foggo, and P. M. Ramsay.
483 2009. Ecology and conservation status of temporary and fluctuating ponds in two areas of southern
484 England. *Aquatic Conservation: Marine and Freshwater Ecosystems* 19: 134- 146

485 Boix, D., Kneitel, J., Robson, B. J., Duchet, C., Zuniga, L., Day, J., Gascon, S., Sala, J., Quintana, X.
486 D. and Blaustein, L. 2016. Invertebrates of Freshwater Temporary Ponds in Mediterranean Climates.
487 In Batzer, D., and Boix, D. 2016 (eds.) *Invertebrates in Freshwater Wetlands*. Springer International
488 Publishing, Switzerland pp. 141

489 Borcard, D., Legendre, P. and Drapeau, P. 1992. Partialling out the spatial component of ecological
490 variation. *Ecology* 73: 1045-1055

491 Bornette, G., C. Amoros, and N. Lamouroux. 1998. Aquatic plant diversity in riverine wetlands: the
492 role of connectivity. *Freshwater Biology* 39: 267-283

493 Bratton, J. H. 1990. Seasonal Pools: an overlooked invertebrate habitat. *British Wildlife* 2: 22-29

494 Brendonck, L., Michels, E., De Meester, L. and Riddoch, B. 2002. Temporary ponds are not 'enemy-
495 free'. *Hydrobiologia* 486: 147-159

496 BRIG. 2008. UK Biodiversity Action Plan Priority Habitat Descriptions; Ponds. pp 1-101
497 http://jncc.defra.gov.uk/PDF/UKBAP_PriorityHabitatDesc-Rev2010.pdf [Last accessed 18/04/2016]

498 Bronmark, C., and L. Hansson. (editors) 2005. *The Biology of Lakes and Ponds*. Oxford University
499 Press, Oxford

500 Buijse, A. D., H. Coops, M. Staras, L. H. Jans, G. J. Van Geest, R. E. Grifts, B. W. Ibelings, W.
501 Oosterberg, and F. C. J. M. Roozen. 2002. Restoration strategies for river floodplains along large
502 lowland rivers in Europe. *Freshwater Biology* 47: 889-907

503 Castella, E., Béguin, O., Besacier-Monbertrand, A. L., Hug Peter, D., Lamouroux, N., Mayor Siméant,
504 H., McCrae, D., Olivier, J. M. and Paillex, A. 2015. Realised and predicted changes in the
505 invertebrate benthos after restoration of connectivity to the floodplain of a large river. *Freshwater*
506 *Biology* 60: 1131-1146.

507 Clarke, K. R., and R. N. Gorley. 2006. *PRIMER v6: User Manual/Tutorial*. PRIMER E-Ltd:
508 Plymouth, UK

509 Collinson, N. H., J. Biggs, A. Corfield, M. J. Hodson, D. Walker, M. Whitfield, and P. Williams.
510 1995. Temporary and permanent ponds: an assessment of the effects of drying out on the conservation
511 value of aquatic macroinvertebrate communities. *Biological Conservation* 74: 125-133

512 Cottenie, K. 2005. Integrating environmental and spatial processes in ecological community dynamics.
513 *Ecology Letters* 8: 1175-1182

514 Cottenie, K. and De Meester, L. 2003. Connectivity and Cladoceran species richness in a
515 metacommunity of shallow lakes. *Freshwater Biology* 48: 823-832

516 Cottenie, K., Michels, E., Nuytten, N. and De Meester, L. 2003. Zooplankton metacommunity
517 structure: regional vs. local processes in highly interconnected ponds. *Ecology* 84: 991-1000

518 Davy-Bowker, J. 2002. A mark and recapture study of water beetles (Coleoptera: Dytiscidae) in a
519 group of semi-permanent and temporary ponds. *Aquatic Ecology* 36: 435-446

520 De Meester, L., Declerck, S., Stoks, R., Louette, G., Van De Meutter, F., De Bie, T., Michels, E. and
521 Brendonck, L. 2005. Ponds and pools as model systems in conservation biology, ecology and
522 evolutionary biology. *Aquatic Conservation: Marine and Freshwater Ecosystems* 15: 715-725

523 Dell, A. I., R. A. Alford, and R.G. Pearson. 2014. Intermittent pool beds are cyclic habitats with
524 distinct wet, moist and dry phases. *Plos One* 9: 1-11

525 Della Bella, V., M. Bazzanti, and F. Chariotti. 2005. Macroinvertebrate diversity and conservation
526 status of Mediterranean ponds in Italy: water permanence and mesohabitat influence. *Aquatic*
527 *Conservation: Marine and Freshwater Ecosystems* 15: 583-600

528 Drake, M. 2001. The importance of temporary waters for Diptera (true-flies). *Freshwater Forum* 17:
529 26-39

530 Dufrêne, M. and P. Legendre. 1997. Species assemblages and indicator species: The need for a
531 flexible asymmetrical approach. *Ecological Monographs* 67: 345-366

532 Florencio, M., C. Díaz-Paniagua, C. Gómez-Rodríguez, and L. Serrano. 2014. Diversity patterns in a
533 macroinvertebrate community of a temporary pond network. *Insect Conservation and Diversity* 7: 4-
534 21

535 Gergel, S. E. 2002. Assessing cumulative impacts of levees and dams on floodplain ponds: a neutral-
536 terrain model approach. *Ecological Applications* 12: 1740-1754

537 Hassall, C., Hollinshead, J. and Hull, A. 2011. Environmental correlates of plant and invertebrate
538 species richness in ponds. *Biodiversity conservation* 20: 3189-3222

539 Hassall, C., Hill, M. Gledhill, D. and Biggs, J. 2016. The ecology and management of urban
540 pondscapes. In: Francis, R., Millington, J. D. A. and Chadwick, M. A. (ed.) 2016. *Urban landscape*
541 *ecology: Science, Policy and Practice*, Routledge:Abingdon, UK.

542 Heino, J., Melo, A. S., Siqueira, T., Soininen, J., Valanko, S. and Bini, L. M. 2014. Metacommunity
543 organisation, spatial extent and dispersal in aquatic systems: patterns processes and prospects.
544 *Freshwater Biology* 60: 845-869

545 Hinden, H., Oertli, B., Menetrey, N., Sager, L. and Lachavanne, J. 2005. Alpine pond biodiversity:
546 what are the related environmental variables. *Aquatic Conservation: Marine and Freshwater*
547 *Ecosystems* 15: 613-624

548 Helfield, J. M., J. Engstrom, J. T. Michel, C. Nilsson, and R. Jansson. 2012. Effects of river
549 restoration on riparian diversity in secondary channels of the Pite River, Sweden. *Environmental*
550 *Management* 49: 130-141

551 Hurlbert, S. H. 1971. The nonconcept of species diversity: a critique and alternative parameters.
552 *Ecology* 52: 577–585.

553 Jeffries, M. J. 2015. Flood, drought and the inter-annual variation to the number and size of ponds and
554 small wetlands in an English lowland landscape over three years of weather extremes. *Hydrobiologia*.
555 768: 255-272.

556 JNCC, 2015. <http://jncc.defra.gov.uk/page-3408> [Last accessed 15/07/2015]

557 Lake, S., N. Bond, and P. Reich. 2006. Floods down rivers: from damaging to replenishing forces.
558 *Advances in Ecological Research* 39: 41-62

559 Legendre, P., and H. J. B. Birks. 2012. From classical to canonical conservation, *in* Birks, H. J. B., A.
560 F. Lotter, S. Juggins, and J. P. Smol. (editors) *Tracking Environmental Change Using Lake*
561 *Sediments, Volume 5: Data Handling and Numerical Techniques*. Springer, Dordrecht, 201-248

562 Legendre, P. and Gallagher, E. D. 2001. Ecologically meaningful transformations for ordination of
563 species data. *Oecologia* 129: 271-280

564 Leibold, M. A., Holyoak, M., Mouquet, N., Amarasekare, P., Chase, J. M., Hoopes, M. F., Holt, R. D.,
565 Shurin, J. B., Law, R., Tilman, D., Loreau, M. and Gonzalez, A. 2004. The metacommunity concept:
566 a framework for multi-scale community ecology. *Ecology Letters* 7: 601-613

567 Lott, D. 2001. Ground beetles and rove beetles associated with temporary ponds in England.
568 *Freshwater Forum* 17: 40-53.

569 Marsh T.J., Parry S., Kendon M.C. & Hannaford J. 2013. The 2010-12 drought and subsequent
570 extensive flooding. Centre for Ecology & Hydrology, Wallingford, U.K. 54 pp.

571 McCabe, D. J. and Gotelli, N. J. 2000. Effects of disturbance frequency, intensity and area on
572 assemblages of stream macroinvertebrates. *Oecologia* 124: 270-279

573 Merritt, R. W. and K. W. Cummins. 1996. An introduction to the aquatic insects of North America,
574 Kendall/Hunt Publishing Company: Debuque, IA

575 Ng, I. S. Y., Carr, C. M. and Cottenie, K. 2009. Hierarchical zooplankton metacommunities;
576 distinguishing between high and limiting dispersal mechanisms. *Hydrobiologia* 619: 133-143

577 Nicolet, P. 2001. Temporary ponds in the UK: a critical diversity resource for freshwater plants and
578 animals. *Freshwater Forum* 17: 16-25

579 Nicolet, P., J. Biggs, G. Fox, M. J. Hodson, C. Reynolds, M. Whitfield, and P. Williams. 2004. The
580 wetland plant and macroinvertebrate assemblages of temporary ponds in England and Wales.
581 *Biological Conservation* 120: 261-278

582 Nilsson, C., C. A. Reidy, M. Dynesius, and C. Revenga. 2005. Fragmentation and flow regulation of
583 the worlds large river systems. *Science* 308: 405-408

584 Paillex, A., E. Castella, P. S. E. Zu Ermgassen, and D. C. Aldridge. 2015. Testing predictions of
585 change in alien and native macroinvertebrate communities and their interaction after the restoration of
586 a large river floodplain. *Freshwater Biology* 60: 1162-1175

587 Paillex, A., S. Doledec, E. Castella, S. Merigoux, and D. Aldridge. 2013. Functional diversity in a
588 large river floodplain: anticipating the response of native and alien macroinvertebrate to the
589 restoration of hydrological connectivity. *Journal of Applied Ecology* 50: 97-106

590 Quinlan, J. R. 1993. Combining instance-based and model-based learning. In *Proceedings of the tenth*
591 *International Conference on Machine Learning*. Morgan Kaufmann: California, USA. pp 236-243

592 Reckendorfer, W., C. Baranyi, A. Funk, and F. Schiemer. 2006. Floodplain restoration by reinforcing
593 hydrological connectivity: expected effects on aquatic mollusc communities. *Journal of Applied*
594 *Ecology* 43: 474-484

595 Savage, A. A. 1989. Adults of the British aquatic Hemiptera Heteroptera: a key with ecological notes.
596 *Freshwater Biological Association Scientific Publication No. 50*, Freshwater Biological Association:
597 Cumbria

598 Shiel, R. J., J. D. Green, and D. L. Nielsen. 1998. Floodplain diversity: why are there so many species?
599 *Hydrobiologia* 387/388: 39-46

600 Starr, S. M., J. P. Benstead, and R. A. Sponseller. 2014. Spatial and temporal organization of
601 macroinvertebrate assemblages in a lowland floodplain ecosystem. *Landscape Ecology* 29: 1017-1031

602 Tachet, H., P. Richoux, M. Bournaud and U. Usseglio-Polatera. 2003. *Invertébrés d'eau douce*
603 *systematique, biologie, ecologie*, CNRS Editions: Paris

604 ter Braak, C. J. F., and P. Šmilaur. 2002. CANOCO reference manual and CanoDraw for Windows
605 users guide: software for canonical community ordination (version 4.5). Microcomputer Power, Ithaca,
606 New York

607 Tockner, K., F. Malard, and J.V. Ward. 2000. An extension of the flood pulse concept. *Hydrological*
608 *Processes* 14: 2861-2883

609 Tockner, K., and J.A. Stanford. 2002. Review of: riverine flood plains: present state and future trends.
610 *Environmental Conservation* 29: 308-330

611 Tockner, K., M. Puschm, D. Borchardt, and M. S. Lorang. 2010. Multiple stressors in coupled river-
612 floodplain ecosystems. *Freshwater Biology* 55: 135-151

613 Van De Meutter, F., De Meester, L. and Stoks, R. 2007. Metacommunity structure of pond
614 macroinvertebrates: effects of dispersal mode and generation time. *Ecology* 88: 1687-1695

615 Vanschoenwinkel, B., C. Vries, M. Seaman, and L. Brendonck. 2007. The role of metacommunity
616 processes in shaping invertebrate rock pool communities along a dispersal gradient. *Oikos* 116: 1255-
617 1266

618 Ward, J. V. K. Tockner, and F. Schiemer. 1999. Diversity of floodplain river ecosystems: ecotones
619 and connectivity. *Regulated Rivers: Research and Management* 15: 125-139

620 Waterkeyn, A., P. Grillas, B. Vanschoenwinkel, and L. Brendonck. 2008. Invertebrate community
621 patterns in Mediterranean temporary wetlands along hydroperiod and salinity gradients. *Freshwater*
622 *Biology* 53: 1808-1822

623 Williams, D. D. 1996. Environmental constraints in temporary freshwaters. *Journal of the North*
624 *American Benthological Society* 15: 634-650

625 Williams, D. D. 1997. Temporary ponds and their invertebrate communities. *Aquatic Conservation:*
626 *Marine and Freshwater Ecosystems* 7: 105-117

627 Williams, D. D. 2006. *The Biology of Temporary Waters*. Oxford University Press, Oxford

628 Williams, P., J. Biggs, G. Fox, P. Nicolet, and M. Whitfield. 2001. History, origins and importance of
629 temporary ponds. *Freshwater Forum* 17: 7-15

630 Williams, P., M. Whitfield, J. Biggs, S. Bray, G. Fox, P. Nicolet, and D. Sear. 2003. Comparative
631 diversity of rivers, streams, ditches and ponds in an agricultural landscape in Southern England.
632 *Biological Conservation* 115: 329-341

633 Williams, P., Whitfield, M. and Biggs, J. 2008. How can we make new ponds biodiverse? A case
634 study monitored over 7 years. *Hydrobiologia* 597, 137-148

635 Witten I.H., Frank, E. and Hall, M, A. 2011. *Data Mining: Practical Machine Learning Tools and*
636 *Techniques*. 3rd ed. Morgan Kaufmann: Burlington, USA

637 Witten, I.H. and Frank. E. 2000. *Data Mining: Practical Machine Learning Tools and Techniques with*
638 *Java Implementations*. Morgan Kaufmann: California, USA.

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647 **Tables**

648 Table 1 - Summary table of measured environmental variables for ephemeral and perennial ponds
 649 across the floodplain meadow sites; SWS: pond surface area shaded, PMS: pond margin shaded, EM:
 650 emergent macrophytes, SM: submerged macrophytes, COND: conductivity, DO: dissolved oxygen.

	Perennial (n = 20)				Ephemeral (n = 14)			
	Mean	Std.Error	Min	Max	Mean	Std.Error	Min	Max
Area (m ²)	828	589	13	11923	230	90	10	1258
Depth (cm)	65	5	27	>100	26	7	8	>100
SWS (%)	9	6	0	93	2.9	2	0	30
PMS (%)	10	5	0	97	7.3	6	0	85
EM (%)	11	3	1	45	37	8	0	87
SM (%)	25	4	4	73	36	9	0	100
pH	8.3	0.1	7.2	9.1	7.5	0.2	6.4	8.7
COND (μS cm ⁻¹)	773	59	422	1494	418	55	80	987
DO (%)	89	4	28	112	78	6	55	120

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665 Table 2 - Summary table (\pm SE) of macroinvertebrate diversity within the ephemeral and perennial
 666 floodplain meadow pond sites. * indicates statistically significant difference ($p < 0.05$) between
 667 ephemeral and perennial ponds.

	Perennial meadow ponds	Ephemeral meadow ponds	All ponds
Total taxon	164	93	173
Mean taxa *	53 (± 2.71)	19 (± 3.21)	39 (± 3.60)
Rarefied taxa richness*	23	14	19
Mean abundance *	3155 (± 292.64)	671 (± 200)	2132 (± 284)
Multivariate dispersion (MVDISP)	0.73	1.564	n/a
Total number of ponds supporting at least one taxa with a conservation designation	8	5	13
Taxa with a conservation designation	<i>Berosus luridus</i> , <i>Ilybius subaeneus</i> , <i>Agabus conspersus</i> , <i>Hygrotus nigrolineatus</i> , <i>Rhantus frontalis</i>	<i>Helophorus dorsalis</i> , <i>Paracymus scutellaris</i> , <i>Hygrotus nigrolineatus</i> , <i>Rhantus frontalis</i>	

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682 Table 3 - Top 6 aquatic macroinvertebrate taxa identified as indicator species for ephemeral or
 683 perennial ponds. * = p<0.05, ** = P<0.01.

Ephemeral ponds	Stat	Perennial ponds	Stat
Collembola**	0.93	Dytiscidae larvae**	0.97
<i>Hesperocorixa sahlberghi</i> *	0.66	<i>Crangonyx pseudogracilis</i> **	0.95
		<i>Stagnicola palustris</i> **	0.95
		Corixidae nymph**	0.92
		Physidae**	0.90
		<i>Radix Balthica</i> **	0.89

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701 **Figure Captions**

702 **Figure 1** Mean percentage of taxonomic orders recorded within the perennial and ephemeral
703 floodplain meadow ponds in this study for selected macroinvertebrate groups.

704 **Figure 2** Macroinvertebrate abundance (a), taxonomic richness (b) and rarefied taxonomic
705 richness (based on 30 individuals drawn randomly from a sample) recorded within
706 different mesohabitat units within perennial and ephemeral ponds. Central black bar =
707 median, box = interquartile range, whiskers = total maximum and minimum range.
708 Open circle = outlier defined on the basis of being >1.5 times the interquartile range
709 from the rest of the values, * = outlier defined on the basis of being >3 times the
710 interquartile range from the rest of the scores.

711 **Figure 3** Proportion (mean %) of functional feeding group (a) and dispersal type (b) among
712 ephemeral and perennial pond communities.

713 **Figure 4** RDA ordination of site plots for perennial and ephemeral floodplain meadow pond
714 Hellinger transformed macroinvertebrate assemblages: (a) site plot with significant
715 environmental parameters shown and (b) taxon richness bubble plot. Empty circles =
716 perennial ponds, filled circles = ephemeral ponds. Note - the size of each bubble is
717 proportional to the absolute taxonomic richness.

718 **Figure 5** The unique and combined influence of physical and chemical, biological and spatial
719 variables on macroinvertebrate composition. Values represent the proportion of the
720 total variation (1.00). Percentage contribution of the total variance is presented in
721 parenthesis.

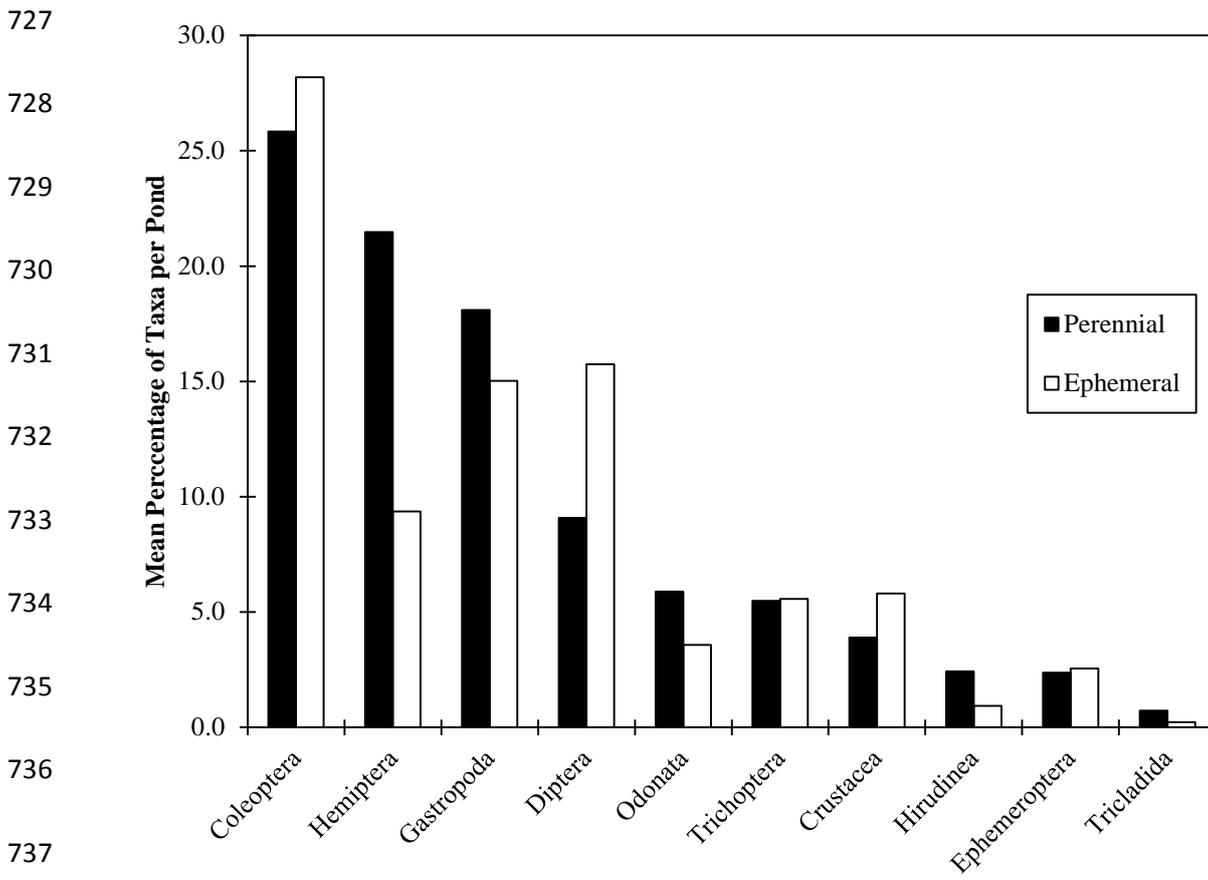
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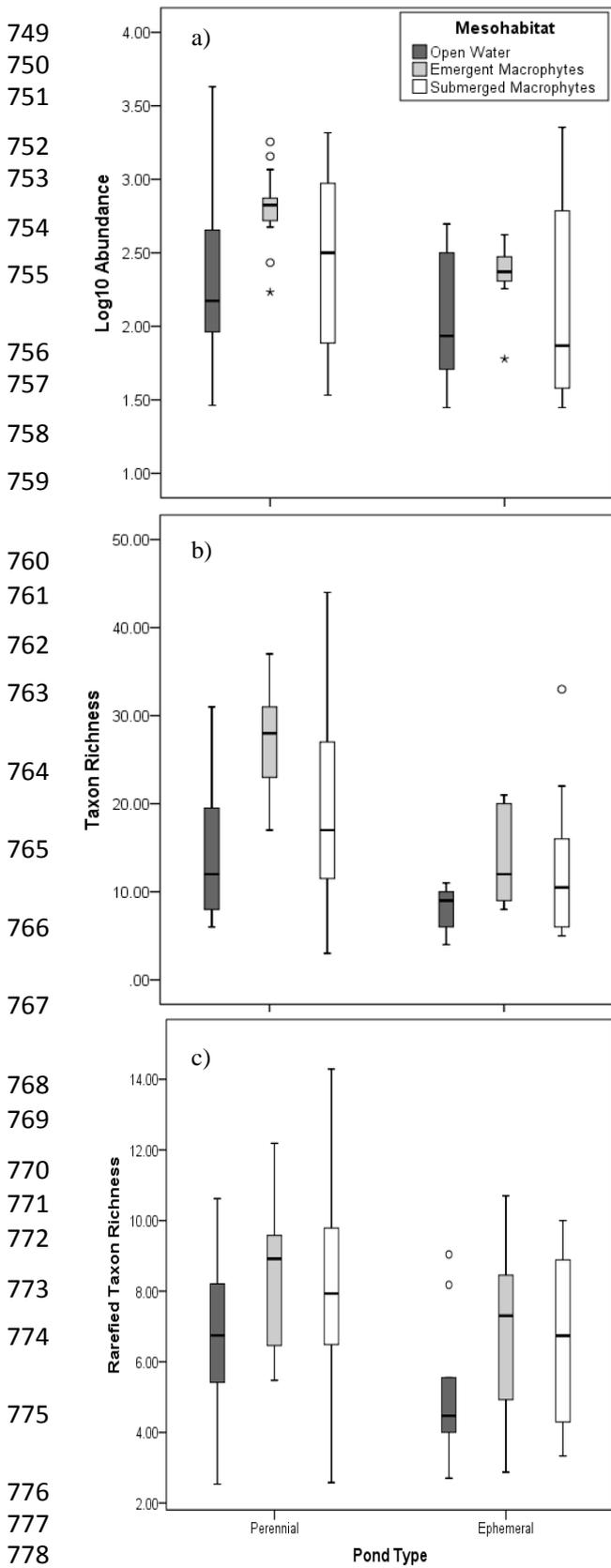
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726 **Figure 1**

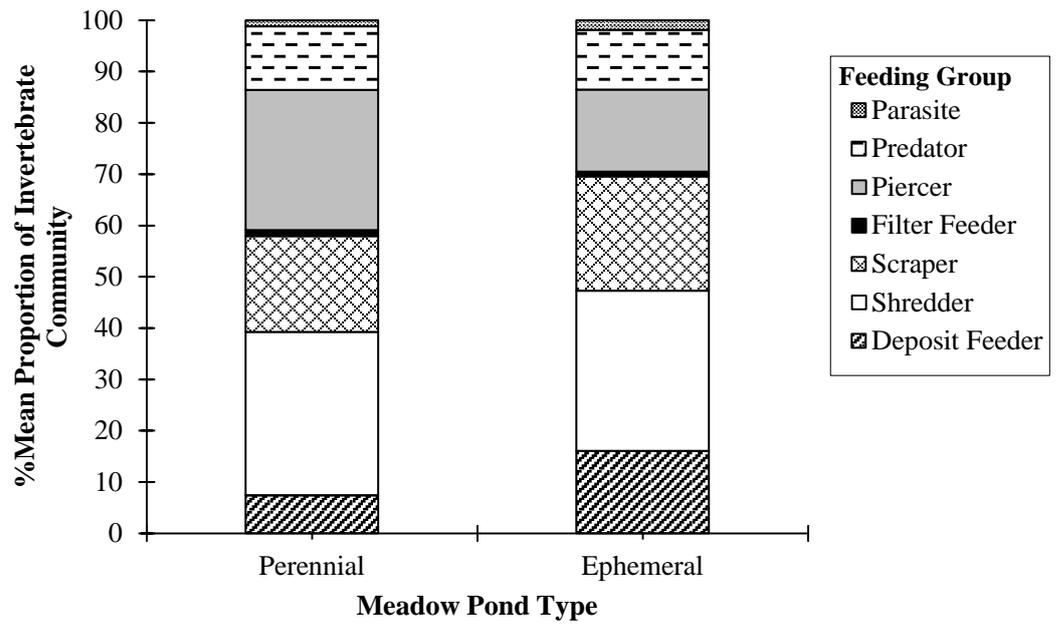


748 **Figure 2**

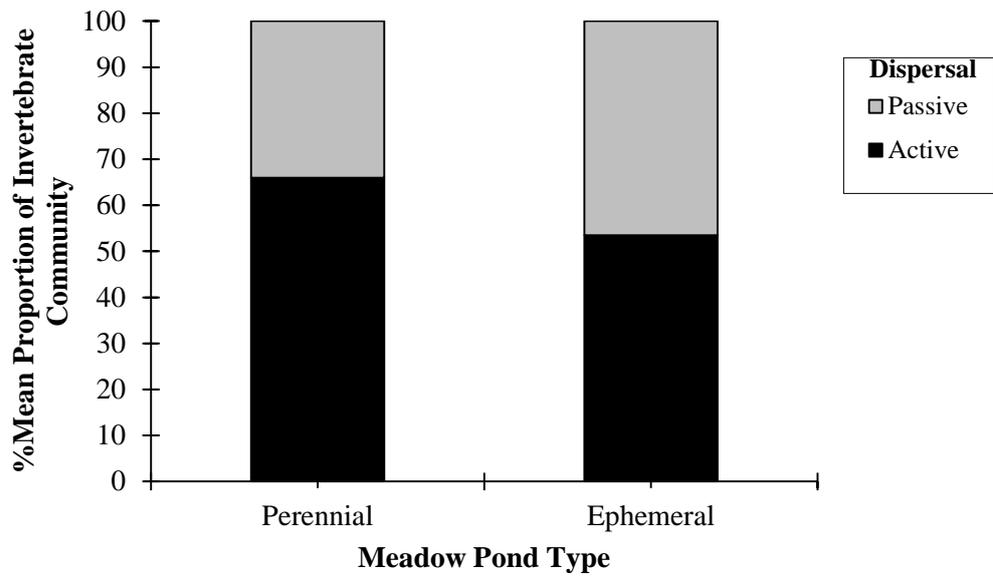


781 **Figure 3**

(a)

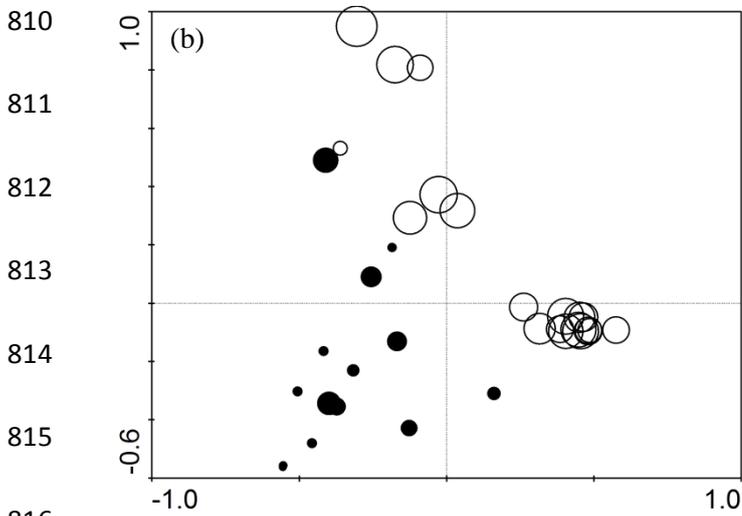
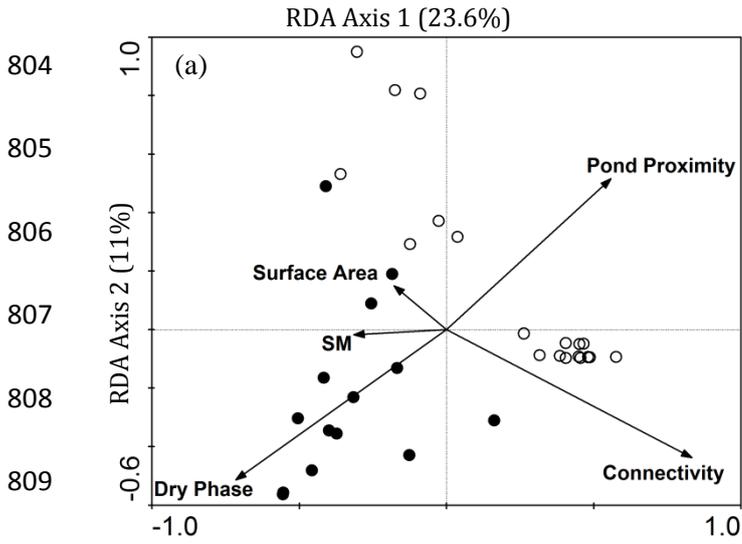


(b)



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803 **Figure 4**



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827 **Figure 5**

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