1	Modelling biodiversity distribution in agricultural landscapes to support ecological
2	network planning
3	
4	Hannah L MOSSMAN*
5	Christopher J PANTER
6	Paul M DOLMAN
7	
8	School of Environmental Sciences, University of East Anglia, Norwich, NR4 7TJ
9	
10	* Current address: Division of Biology and Conservation Ecology, School of Science and the
11	Environment, Manchester Metropolitan University, Chester St, Manchester, M1 5GD.
12	
13	Corresponding Author:
14	Hannah L Mossman, Division of Biology and Conservation Ecology, School of Science and
15	the Environment, Manchester Metropolitan University, Chester St, Manchester, M1 5GD.
16	Tel: +44 (0) 161 247 1186. Email: <u>h.mossman@mmu.ac.uk</u>
17	
18	Co-author email addresses:
19	Christopher J Panter: c.panter@footprint-ecology.co.uk
20	Paul M Dolman: p.dolman@uea.ac.uk

22 Abstract

23 Strategic approaches to biodiversity conservation increasingly emphasise the restoration of 24 ecological connectivity at landscape scales. However, understanding where these connecting 25 elements should be placed in the landscape is critical if they are to provide both value for 26 money and for biodiversity. For such planning to be effective, it is necessary to have 27 information of the distributions of multiple taxa, however, this is of poor quality for many 28 taxa. We show that sparse, non-systematically collected biological records can be modelled 29 using readily available environmental variables to meaningfully predict potential biodiversity 30 richness, including rare and threatened species, across a landscape. Using a large database of 31 ad-hoc biological records (50 501 records of 502 species) we modelled the richness of 32 wetland biodiversity across the Fens, a formerly extensive wetland, now agricultural 33 landscape in eastern England. We used these models to predict those parts of the agricultural 34 ditch network of greatest potential conservation value and compared this to current strategic 35 network planning. Odonata distribution differed to that of other groups, indicating that single taxon groups may not be effective proxies for other priority biodiversity. Our results 36 37 challenged previous assumptions that river channels should comprise the main connecting elements in the Fens region. Rather, areas of high ditch density close to a main river are 38 39 likely to be of greater value and should be targeted for enhancement. This approach can be 40 adopted elsewhere in order to improve the evidence-base for strategic networks plans, 41 increasing their value for money. 42

43 HIGHLIGHTS

• We used ad-hoc biological data to model landscape-scale wetland species richness.

- Models were used to assess and improve a proposed ecological connectivity network.
- Our evidence-based network was shorter and connected areas of higher richness.
- Our results challenge previous assumptions of important network elements.
- Odonata were poor proxies for other groups of wetland species.
- 49

50 1. Introduction

51

52 Habitats are increasingly fragmented. Furthermore, in human landscapes, habitat patches are

53 often surrounded by land uses that are potentially hostile to dispersal, increasing functional

- isolation (Nowicki et al., 2014). Such habitat fragmentation and isolation increase local
- 55 population vulnerability to extinction and reduced dispersal opportunities limit species'

56 ability to respond to climate change, further reducing biodiversity resilience (Hill et al., 57 2002). Strategic approaches to conservation are, therefore, increasingly focused at the 58 restoration of landscape connectivity by the creation of movement corridors, stepping stones 59 or by improving landscape permeability (Dolman, 2012; Lawson et al., 2012; Saura et al., 60 2014). However, the nature, size and placement of these connecting elements are critical if 61 investment of finite funds and land resources are to give optimal returns. There are several 62 key issues to the success of landscape connectivity; identifying what species should be targeted within a landscape (Dolman et al., 2012), ensuring that the connectivity elements 63 64 comprise habitats that suit these species and establishing where these connecting elements 65 should be placed.

66

67 Ecological networks are often designed to enhance the metapopulation viability of individual 68 high profile species, such as top predators (Klar et al., 2012) or other mobile species (Bani et 69 al., 2002). However, the ability of such species to act as connectivity umbrellas for assemblages of other species may be limited (Cushman & Landguth, 2012) because the 70 71 suitability of the habitat and type of connecting element differs amongst taxa. For example, 72 while linear field margins may provide connectivity to some generalist butterflies (Delattre et 73 al., 2010), they may act as sinks to other taxa (Krewenka et al., 2011). Similarly, hedgerows 74 are often purported to provide suitable corridors for woodland species, but may only provide 75 habitat for woodland edge species (Liira & Paal, 2013). The planning of landscapes to 76 provide resilience for assemblages of regional biodiversity therefore requires the 77 consideration of multiple relevant taxa (Zulka et al., 2014).

78

79 Decisions regarding the optimum placement of connecting elements should be made using 80 evidence of the current and potential distribution of a full complement of target species. 81 Existing protected sites that retain a concentration of rare species generally form the focus of 82 connectivity networks (Beier et al., 2011) and the existence of species within these fragments 83 is often well known. However, our understanding of the distribution of species throughout the rest of the landscape is incomplete, with some locations receiving high levels of recording 84 effort and others very little. Poorly recorded areas that are nevertheless potentially suitable 85 86 for a species may harbour unrecorded residual populations, or be more likely to be colonised 87 if both habitat quality and connectivity are improved (Lawson et al., 2014). Unsystematically 88 collected biological data therefore do not provide a reliable assessment of conservation value 89 or potential across a landscape. This results in reliance on expert opinion in the design of

landscape connectivity (Beier et al., 2009; Eycott et al., 2011). However, if the patchy nature
of recording effort is accounted for in the analysis (Kéry, 2011), ad hoc biological data can be
exploited to provide more objective design of landscape connectivity.

93

94 In this study, we use the Fens, a formerly extensive wetland system in eastern England, to 95 demonstrate how connectivity planning can be informed by modelling ad hoc biological 96 records with easily obtainable, landscape-scale environmental data. Remaining wetland 97 habitat in the Fens is highly fragmented and isolated within an intensive agricultural 98 landscape, but there is a high potential for connectivity through enhancing management of 99 linear drainage ditches. Ditches in intense agricultural areas are often rather different to natural streams (Herzon & Helenius, 2008), supporting lower biodiversity (Williams et al., 100 101 2004); however, they can act as reservoirs for important regional wetland biodiversity (Simon & Travis, 2011). Biological recording within the wider Fens landscape is extremely sparse, so 102 simple mapped biological richness cannot be used as an evidence base for selecting 103 104 potentially biodiverse ditches for improved management or in the design of connectivity networks. Recent attempts at strategic planning (e.g. Fens for the Future Partnership, 2012) 105 106 have therefore relied on a combination of expert opinion and untested assumptions of where 107 this targeted management should be placed.

108

109 We take the approach of modelling potential biodiversity value in relation to underlying environmental factors and landscape context, to predict where in the landscape targeted 110 111 management to enhance habitat quality will have greatest potential to support biodiversity and enhance connectivity. We use an extensive but unevenly distributed database of 67,395 112 113 ad hoc biological records to model the richness of groups of wetland species across the Fens 114 landscape in relation to a range of coarse-scale environmental and landscape factors. Using 115 these models, we aim to: 1) predict and map the potential richness of groups of wetland 116 species in order to identify parts of the landscape of greatest potential conservation value; 2) 117 apply these maps of predicted biodiversity potential to assess current strategic planning maps.

118

119 **2.** Methodology

120 *2.1 Study area*

121 The Fens, covering almost 4,000 km² of eastern England (Fig. 1), was formerly an extensive 122 wetland area but only 1% of wetland habitat remains. This habitat is concentrated in six key 123 protected areas, which are each small (mean area 819 ha) and isolated within the country's

- most important arable agricultural landscape. More than 20 million km of ditches and
- drainage channels criss-cross the Fens landscape and by targeting selected ditches for
- 126 enhanced management, the ditch network presents an excellent opportunity for increasing
- 127 both habitat area and connectivity for wetland species. However, the current conservation
- 128 value of large parts of this landscape is poorly known.
- 129

130 *2.2 Biological data*

All available species observations (records) were collated for the period of 1987-2012 from 131 132 the 4147 1-km squares wholly or partly within the Fens Natural Character Area boundary 133 (Natural England, 2013), with an extension (3 km from the boundary) to include Chippenham 134 Fen, one of the three important relict fen sites in the Fens. Records were compiled from Local Biological Records Centres, the National Biodiversity Network (NBN) gateway, national and 135 136 county natural history and recording societies whose records were not available via NBN, and 137 unpublished documents or reports. Records sent to Biological Records Centres and societies are validated by expert county recorders. Although NBN data may include some unvalidated 138 records submitted by the public, our collated species lists were validated by a range of local 139 140 taxonomic experts. Records were managed using RECORDER 6 software (Joint Committee 141 for Nature Conservation, Peterborough, UK). The study period (1987-2012 inclusive) was selected as a compromise between reflecting the current or recent distribution of wetland 142 143 species and including sufficient records in the dataset to capture rare species and the potential distribution of sparsely recorded taxonomic groups. There may have been local extinctions 144 145 since 1987 due to local changes in habitat quality, nevertheless the landscape predictors we consider will indicate the biodiversity potential should habitat and connectivity be restored. 146 147 The majority (74%) of records were resolved to a spatial resolution of 1 km or finer and these 148 were aggregated and analysed at the scale of 1-km grid cells. Tetrad records were assigned to 149 all of the four 1-km squares comprising the tetrad; species records at coarser spatial 150 resolutions were excluded. A small number of records of taxa not recorded to species level were removed. Records of marine species were excluded, but those tolerant of brackish 151 152 conditions were retained. Following additional filters described below (e.g. removal of 153 coastal squares), a database of 255 291 records remained, of which 50 501 were records of 154 wetland plants (including conservation priority species) and conservation priority wetland 155 invertebrates. Conservation priority species were recognised as those designated as UK 156 Biodiversity Action Plan, Global and UK Red Data Book (except Least Concern), Nationally Rare, Nationally Scarce or Nationally Notable A and B, according to JNCC (2012), plus 157

undesignated species with >25% of their UK distribution occurring in the Fens region –
hereafter referred to as 'Fens Specialists'.

160

161 The richness of groups of wetland species were used as the biological response variables. 162 Seven widely recorded groups of wetland species were selected for modelling that were 163 considered good indicators of ditch quality: all Odonata (dragonflies and damselflies, 28 164 species), wetland plants (212 taxa), fully aquatic plants (137 taxa) and conservation priority species (including plants and invertebrates) dependent on aquatic (fully aquatic and 165 166 submerged aquatic habitats, 90 species), littoral (aquatic margins, 109 species) or wetland 167 (208 species) habitats, and Fen Specialists (58 species). Wetland plant species were defined 168 as all Characeae (stoneworts, multi-cellular branched macro-algae) and those vascular plant species associated with freshwater (aquatic, wetland or seasonally wet) habitats selected from 169 170 Hill et al. (2004) with Ellenberg moisture values \geq 7 (species with Ellenberg salinity values 171 of >5 were excluded). Aquatic plants were a sub-set of the wetland plants, classified with reference to existing lists by Palmer et al. (2013) and Mountford and Arnold (2006). The 172 autoecological requirements of conservation priority species and their association with 173 174 wetland, aquatic and littoral habitats, were classified following Mossman et al. (2012) and 175 Dolman et al. (2012).

176

177 2.3 Environmental predictors

The aim of this analysis was to predict the distribution of wetland species across the drainage 178 179 ditch network of the arable landscape based on readily-available, coarse-scale environmental variables. Wetland Sites of Special Scientific Interest (SSSIs) were considered to be 180 181 reservoirs and potential sources of high quality biodiversity, therefore 1-km squares including 182 any part of a wetland SSSI were excluded from modelling. Wetland SSSIs were identified 183 based on the SSSI citation description (available at www.sssi.naturalengland.org.uk), with 184 wetland habitats considered to include ponds, gravel pits, wet woodland or carr, fen, bog, 185 grazing marsh and wet common.

186

187 Seventeen environmental predictors were initially selected as candidates for modelling (Table

188 1) based on ready availability across the study landscape and considered, *a priori*, to

189 potentially influence ditch biodiversity. A single value of each variable was calculated for

190 each 1 km square. The mean elevation above sea level, presence of an A or B road and the

191 distance from the centre of each 1-km square to the nearest wetland SSSI, Fenland island and

the edge of the Fen basin were calculated. Previous work has suggested that ditches with

highest conservation value are located near to the edge of the Fen basin or close to Fen

islands (Mountford & Arnold, 2006); the reasons for this are unclear, but may relate to high

195 water quality. Fen islands were delimited as areas of >0.1 km² with an elevation of \ge 5 m, and

the Fen basin defined as the 5 m contour boundary.

197

198 The soils of the Fens area are dominated by silt and peaty soil types. The percentages of each 199 1-km square comprising silt and selected peat soil types (Table 1) were calculated. Ditch 200 isolation from main channels and from tidal influence were considered potentially important 201 determinants of water quality, saline influence and thus of biodiversity richness. We 202 calculated the shortest network distance along the ditches and rivers network (extracted from 203 the Ordnance Survey (OS) surface water polylines, converted into a raster of 35m cells), 204 from the centre of each 1-km square to the nearest main channel/river and to the tidal 205 boundary, calculated in ArcGIS Spatial Analyst tools. Network distances were not weighted by ditch size or type, such that all cells were assigned a value of 1. A cell size of 35 m was 206 sufficient to connect any small breaks in the polylines due to mapping error or underground 207 208 drains, but was considered small enough to prevent falsely connecting ditches in close 209 proximity that are not connected through surface water drainage. Some manual connections 210 were imposed on the network due to large breaks in the mapped surface, for example due to 211 bridges or pumping stations. Ditch density in each 1-km was calculated from OS polylines, which defines both banks of ditches wider than 2 m; since ditches of <2 m in width are only 212 213 defined with one polyline, ditch density is an index that reflect both linear length and ditch 214 area.

215

216 The grades of the Agriculture Land Classification were used as proxies for potential 217 agricultural productivity, land-use intensity and therefore quality of both water and banksides; this is an ordinal scale (1-5) where grade 1 is best agricultural land. The combined 218 219 percentage cover of grades 3 and 4, comprising the lowest quality agricultural land and 220 therefore representing the lowest intensity of agricultural land-use (no land was classified as grade 5 in our study region), was used as a candidate predictor. The dominant land use in the 221 222 Fens region is arable; the percentage of each 1-km square comprising un-intensively managed 223 grassland (defined from Land Cover Map (Morton et al., 2011)) classes of Rough/Neutral 224 Grassland) was therefore considered of interest. The percentage of urban land use was also 225 calculated from OS data.

226

- 227 Inter-correlation among predictor variables was investigated using Pearson's correlation
- 228 coefficient and considered large enough to potentially have an effect on the models if r > 0.5,
- following Freckleton (2002). Distance to the Fen basin was strongly correlated with distance
- to the nearest wetland SSSI (r=0.533), network distance to the tidal boundary (r=-0.523) and
- percentage of silt soils (r=0.536). Distance to the Fen basin was therefore excluded from the
- 232 modelling, whilst the other variables were retained.
- 233

234 Due to comprehensive county flora, plant species recording effort was substantially greater in

- 235 Norfolk and Suffolk relative to other counties. Therefore, to avoid spurious identification of
- any environmental factor that differed between these and other counties, when modelling the
- response of wetland and aquatic plant variables to environmental and landscape context
- indicators, we included the two county groups as a binary covariate (0 = no flora, 1 = flora).
- 239

A number of 1-km squares were excluded from the models because they contained no surface
water, the surface water was more than 70 m from the nearest surface water feature (thus
indicating the feature was likely to be a pond rather than a ditch, contained part of a wetland
SSSI, or comprised >50% coastal area (defined using the Wash SSSI). This resulted in 3,745
1-km squares being used in analyses.

245

246 2.4 Model construction

247 2.4.1 Accounting for recording effort

It is well known that not all species present at a site will be detected and that this poses 248 249 challenges for analysis (Chen et al., 2013), as species richness is underestimated and 250 coefficients with environmental variables are closer to zero. Spatial variation in recorder 251 effort can have severe consequences for models, as environmental variables that are 252 correlated with recording effort may be spuriously identified as being related to species 253 richness. Hierarchical occupancy modelling can address these problems by utilising repeated 254 visits to the same site to estimate detection probabilities (MacKenzie & Kendall, 2002) and 255 thus has applications for analysing citizen science data (Isaac et al., 2014). Despite extensions 256 to deal with multiple species (Dorazio & Royle, 2005), application to datasets such as ours is 257 challenged by, for example, uncertainty in defining what represents a discrete 'visit', and 258 absence of information on visits that did not contribute species records to the data. An 259 alternative approach to addressing spatial variation in recorder effort is to include a proxy for

260 recorder effort as a covariate (Hill, 2011), allowing the conditional effects of environmental variables on species richness to be assessed while controlling for recorder effort. We use the 261 262 total number of records in a 1-km square (i.e. including non-wetland species) as a proxy for 263 recording effort. We expect this relationship to be saturating as species accumulation curves 264 tend to saturate at high numbers of species, so we explored models using either square root 265 number of records or a polynomial term for number of records, using the former as it 266 explained more deviance. Although our method accounts for spatial variation in recorder 267 effort, we are unable to estimate the probability of not detecting a species, so our estimates of 268 species richness should be taken as an index of relative richness.

269

270 2.4.2 Predicting species richness

271 Statistical analyses were performed using the computing environment R (R Core Team, 272 2012). Predictor variables were standardised prior to modelling, with the exception of the 273 number of records. For each response variable, we fitted generalised linear models, with a quasi-poisson error structure to deal with over-dispersion, containing all 16 predictor 274 variables (17 for wetland and aquatic plants owing to the inclusion of county). The full model 275 276 was simplified by backward elimination, judging variable retention by the t-test of β 277 estimates, with a threshold of $\alpha < 0.05$. The resulting minimum models were used to predict 278 the richness of each of the seven wetland species groups in each 1-km square of the study 279 area, with recording effort standardised as the overall median (41 records per 1-km square). 280 For the wetland and aquatic plant response variables, we standardised for the presence of a 281 recent flora by setting the value for all squares as 1.

282

Following Legendre and Legendre (2012), we used variance partitioning to calculate the proportion of total variation in species richness explained by recording effort (total number of records) and by environmental variables. To do this, we constructed models including 1) only environmental conditions, 2) only recording effort and 3) both environmental conditions and recording effort.

288

289 2.5 Comparison of predicted biodiversity richness to the current strategic planning maps

290 The 1-km squares were ranked by the predicted species richness for each of the seven

291 biological response variables separately, where a high rank (low number) was given to

squares with high predicted biodiversity. The mean of these ranks was calculated and

293 mapped. The resulting map of predicted biodiversity was compared to the Fens for the Future

294 Partnership (FFFP) (2012) strategic connectivity plan. The strategic connectivity network consisted of three types of corridors: primary, secondary and landscape (Fens for the Future 295 296 Partnership, 2012). The primary corridor was the priority corridor and aimed to connect three 297 core areas thought to have high biodiversity value, the southern Fens and Ouse Washes, 298 Holme and Woodwalton fens (and associated Great Fens Project restoration area of the 299 Wildlife Trusts), and the Nene Washes. Secondary and landscape corridors aimed to provide 300 additional landscape connectivity; for the purposes of this study, secondary and landscape 301 corridors were combined.

302

303 We designed a new connectivity network that met with the objectives of the strategic

304 connectivity network and the following criteria. Corridors must connect areas of known high

305 biodiversity richness (wetland SSSIs) and presumed high richness, defined as those wetland

Local Wildlife Sites (LWS) that were ≥ 0.25 km² and occurred in areas of high predicted

307 biodiversity (richest \geq 50% of 1-km squares). A single primary corridor was placed to connect

the three core sites identified by the FFFP (2012). All corridors must join to form a

309 continuous network across the region and, where possible, achieve such connectivity by

- 310 passing through areas of greater predicted biodiversity.
- 311

The potential conservation effectiveness of the original strategic plan was compared to that of the corridor network we proposed on the basis of the predicted distribution of wetland biodiversity richness. These were assessed for each corridor strata (primary, secondary) in terms of the length within each quartile of predicted species richness (for each 1-km square, the mean of ranked richness across all the seven species groups). Proposed networks were deemed to be more effective if a greater proportion of the corridors lay within the quartiles predicted to the be most species-rich.

319

320 3. Results

321 *3.1 Effect of the environment on wetland biodiversity richness*

322 Overall, the minimum models explained 27.2 - 63.9% (mean = 40.3%) of the variation in

323 species richness of the seven groups (Table 2), performing best in predicting the richness of

wetland plants and aquatic plants (63.9% and 59.8%, respectively). A substantial part of the

- explained variance was attributed to the independent effect of recorder effort (27.3 76.2%).
- However, 17.1 52.8% of explained variance was attributed to the independent effect of
- environmental variables, and a further 2.5-26.4 % to the joint effect of recorder effort and

environmental variables (Table 2). Species groups with the highest proportion of variance
explained by the environmental variables were Odonata, aquatic species and littoral species
(53%, 36% and 33% respectively).

331

The effects of many environmental predictors were consistent among species groups. Mean elevation above sea level and percentage of urban area were not significant predictors of the richness of any group (Fig. 2). A greater percentage of silt soil was negatively related to species richness of all groups, compared to all types of peat soil (Fig. 2), although the richness of wetland and littoral species were also lower with a greater percentage of deep sand over peat or peat (Fig. 2).

338

339 Richness of all groups, except Fen Specialists, was greater closer to existing wetland SSSIs. The richness of all groups except wetland plants, increased significantly with increasing 340 values of the index of ditch density (Fig. 2). The percentage of grade 3 and 4 agricultural land 341 (i.e. lower land-use intensity) was positively related to richness of Odonata, but not 342 significantly related to the richness of other groups. The richness of Odonata also increased 343 further from the tidal boundary; in contrast, the richness of aquatic species, and wetland and 344 345 aquatic plants was higher closer to the tidal boundary. The distance to a main river was not significantly related to the richness of Fen Specialists and littoral species. Richness of the 346 347 remaining groups was highest closer to a main river, although predicted richness generally decreased when main rivers were located on silt soils or were further from a wetland SSSI 348 349 (Fig. 3). The predicted richness of all groups was low around the coast (Fig. 3).

350

351 *3.2 Biodiversity potential of the proposed network corridors*

352 The combined predicted richness of ditch species suggests that the corridors of the proposed 353 strategic network are generally well placed (Fig. 4, 5). However, comparison of the strategic 354 map and the predicted biodiversity richness indicated that proposed corridors do pass 355 through some areas of lower biodiversity potential (Fig. 4). In contrast, our suggested map 356 achieved a greater proportion of connectivity in areas of high predicted richness (88% of our 357 corridors were located in the richest 50% of squares, compared to 66% of the FFTF corridors) 358 for a shorter overall length (27% shorter, combined primary and secondary corridors) (Fig. 359 5).

360

361 4. Discussion

362 Landscape connectivity and conservation plans are often developed with a reliance on environmental and land cover data (Brooks et al., 2004a), but such broad data can be poor 363 364 surrogates for biodiversity (Araujo et al., 2001; Schindler et al., 2013), particularly for rare or 365 specialist species (Lombard et al., 2003). Ecological planning should consider the identity, 366 distribution and requirements of target species in that region, rather than being based on 367 untested assumptions of where species occur (Brooks et al., 2004b) as such assumptions can 368 lead to inappropriate selection of habitat type or placement of the connecting elements. For 369 example, the previous landscape connectivity plan in the Fens that was based on expert 370 opinion selected the main river channels as a key connecting component (FFTP 2012). Whilst we found that species richness was higher closer to main river channels, rivers 371 372 flowing through areas of silt soils had particularly low predicted species richness, so 373 improvements to management or connectivity in these areas may have limited benefits for 374 wetland biodiversity. This has important implications for other landscapes where as single 375 land cover variable has been the focus of network planning, because without validating with 376 biological data the use of single features can prevent selection of optimal connectivity. 377

378 Increasing ditch density was a significant predictor of species richness for all groups, except 379 wetland plants. The ditch density was a particularly strong predictor of priority species (those 380 with a conservation designation) associated with littoral margins. Littoral species are of 381 particular conservation importance in the Fens region, but are often overlooked by conservation interventions compared to submerged aquatic species (Mossman et al. 2012). 382 383 Thus specifically targeting areas of high ditch density close to rivers for improved management, rather than the main river channels themselves, would substantially add 384 385 conservation value. This highlights the importance of considering the identity and 386 requirements of the species that are the priorities for conservation and connectivity in a 387 region or a landscape.

388

Several broad and readily available landscape variables, such as distance to a protected site (SSSI) and cover of silt soils, were important predictors of biodiversity. Thus, such variables can be used to select areas for restoration or connectivity. The consistent negative response of species richness to silt soils may be related to reduced water quality, since sediment nutrient concentrations are higher in finer particle soils (Ockenden et al., 2012), or may reflect the contrasting deposition and landuse histories, with peat soils indicating the historic extent of freshwater marshes and earlier reclamation compared to the marine or riverine deposition of silts that were reclaimed for agriculture more recently. Previous studies have found peat
substrates to have distinct flora (Mountford & Arnold, 2006) and support rare invertebrate
species (Foster et al., 1989); the richess of species groups in this study were not strongly
correlated with peat substrates.

400

401 Environmental factors, such as water quality (Twisk et al., 2000) and flow rate (Leslie et al., 402 2012), and ditch management type and frequency (Milsom et al., 2004), are known to be important determinants of ditch biodiversity. The inclusion of such variables would certainly 403 404 improve the predictive power of our models. However, such data were not available at 405 suitable resolution across our study area, and the case is likely to be the same in other 406 regions. We suggest that our predictive modelling approach is used in other regions to predict 407 areas of high potential biodiversity value. Following this, the collation or collection of 408 detailed environmental or habitat data may assist the selection of specific sites for 409 management interventions (such as dredging and cutting), within those areas highlighted by 410 the predictive mapping.

411

412 The effects of many environmental predictors were remarkably consistent among species 413 groups. For example, the richness of all groups was significantly greater closer to existing 414 wetland SSSIs. This may be because the high quality SSSI sites have acted as reservoirs of 415 wetland species, although there may be other conditions not included in this study (e.g. water quality) that are also correlated with the distance to the SSSIs. Whilst the responses of most 416 417 groups were consistent, the richness of Odonata increased further from the tidal boundary; in contrast, the richness of aquatic species, and wetland and aquatic plants was higher closer to 418 419 the tidal boundary. This is an important contrast, such that network planning must either take 420 a mixed approach, or select to prioritise either Odonata or remaining groups. Similarly, the 421 value of wooded connectivity networks is rather different for birds, bats and beetles 422 (Boughey et al., 2011; Davies & Pullin, 2007). This adds to previously stated concerns over 423 the use of single taxonomic groups as proxies for other biodiversity (Noss, 1990). Recent 424 work has demonstrated that the addition of habitat characteristics to multi-taxa proxy groups 425 substantially improves the performance of biodiversity surrogates in spatial planning (Di 426 Minin & Moilanen, 2014).

427

Biological records can be modelled with environmental variables to predict biodiversity
richness across landscapes and such models have been widely used to link species

430 distributions from atlas data to land cover data (e.g. Atauri & de Lucio, 2001; Virkkala et al., 2005). Their use here to model species richness of priority biodiversity across multiple taxa 431 432 in the Fens allowed previously held assumptions about the importance of landscape features 433 to be tested. However, the use of such methods has been limited by the lack of detailed atlas 434 data for many taxa in many regions, with data for rare and threatened species and for poorly 435 recorded taxonomic groups (i.e. other than vascular plants, butterflies and odonatan) 436 particularly limited. We show that this problem can be overcome by modelling groups of 437 priority taxa with shared ecological requirements, which allowed us to include species that 438 would be too rare and/or sparsely recorded to model individually. This addresses a significant 439 gap in previous large-scale studies that have omitted due to insufficient data, the rare species 440 that are intended to benefit from the conservation measures. Our approach could be applied to 441 any region or landscape where there has been widespread, albeit patchy biological recording. 442

We were then able to predict potential species richness, including multi-taxa groups of 443 priority species, at a landscape scale and used the model predictions to make an evidence-444 445 based landscape connectivity plan, an improvement on previous plans based on untested 446 expert judgement. Our models predict areas that have the potential for high biodiversity 447 richness, based on their soil and other landscape variables, and we have linked these together 448 with our proposed corridors. However, we do not know if the cells of our predicted corridor 449 currently realise that biodiversity potential with their existing habitat, which could still be improved through enlargement or management, or if they currently have low habitat 450 451 suitability despite high potential on the basis their landscape variables. However, in either 452 case, we predict the potential to enhance biodiversity value and connectivity of those areas to 453 be greater than in areas with lower intrinsic potential and thus we are recommending these 454 areas should be targeted for enhancement.

455

456 The previous attempt to map a strategic connectivity network in the Fens (FFTP 2012) largely concurred with areas of high predicted biodiversity richness. However, our evidence-457 458 based map connected a greater proportion of areas with higher potential for biodiversity 459 richness (22% more of our corridors were located in areas of the highest potential richness) 460 and for a shorter overall length. Targeting areas of higher potential richness over a shorter connectivity length is more cost-effective, allowing remaining funding to be targeted to 461 462 habitat management, a key influence on ditch biodiversity (Milsom et al., 2004). For example, our evidence-based predictive map provides confidence in the strategic targeting of 463

agri-environmental measures and other means to enhance ditch management to those areas of
the wider agricultural landscape that have greatest biodiversity potential for aquatic and
wetland species.

467

468 Evidence-based predictive models, such as those in this study, could also be further 469 developed to inform optimal connectivity plans. For example, predicted potential species 470 richness can be used as a cost surface for circuit theory and other graph theory based models (Galpern et al., 2011; Rayfield et al., 2011). Although we note that the practical realization of 471 any connectivity plan (subjective or objective) will be dependent on opportunity, landowner 472 and other stakeholder interest, and cost (Bergsten and Zetterberg, 2013), it is crucially 473 important to start negotiations based on evidence. Our methodology utilises ad-hoc records, 474 and thus could be applied in any landscape or region where biological records are available, 475 to provide an evidence-base for network planning, including rare species for which 476

- 477 conservation actions are most needed.
- 478

479 Acknowledgements

This work is builds on a project funded by the Environment Agency and supported by the 480 Fens for the Future Partnership. We gratefully acknowledge the support and assistance of 481 Natural England, Environment Agency, National Trust, Royal Society for the Protection of 482 Birds, Lincolnshire Wildlife Trust, Bedfordshire, Cambridgeshire and Northamptonshire 483 Wildlife Trust, Cambridge and Peterborough Environmental Records Centre, Suffolk 484 485 Biological Records Centre, Norfolk Biodiversity Information Service and the Norfolk Biodiversity Partnership. This work would not be possible without the invaluable 486 487 contributions of hundreds of biological recorders and taxonomic experts. We also acknowledge the statistical advice of Martin Sullivan, and the helpful comments of three 488 489 anonymous reviewers.

References

- Araujo, M. B., Humphries, C. J., Densham, P. J., Lampinen, R., Hagemeijer, W. J. M., Mitchell-Jones, A. J. & Gasc, J. P. (2001). Would environmental diversity be a good surrogate for species diversity?, *Ecography*, 24(1), 103-110.
- Atauri, J. A. & de Lucio, J. V. (2001). The role of landscape structure in species richness distribution of birds, amphibians, reptiles and lepidopterans in Mediterranean landscapes, *Landscape Ecology*, 16(2), 147-159.
- Bani, L., Baietto, M., Bottoni, L. & Massa, R. (2002). The use of focal species in designing a habitat network for a lowland area of Lombardy, Italy, *Conservation Biology*, *16*(3), 826-831.
- Beier, P., Majka, D. R. & Newell, S. L. (2009). Uncertainty analysis of least-cost modeling for designing wildlife linkages, *Ecological Applications*, 19(8), 2067-2077.
- Beier, P., Spencer, W., Baldwin, R. F. & McRae, B. H. (2011). Toward best practices for developing regional connectivity maps, *Conservation Biology*, 25(5), 879-892.
- Bergsten, A. & Zetterberg, A. (2013). To model the landscape as a network: A practitioner's perspective, *Landscape and Urban Planning*, *119*, 35-43.
- Boughey, K. L., Lake, I. R., Haysom, K. A. & Dolman, P. M. (2011). Improving the biodiversity benefits of hedgerows: How physical characteristics and the proximity of foraging habitat affect the use of linear features by bats, *Biological Conservation*, 144(6), 1790-1798.
- Brooks, T., da Fonseca, G. A. B. & Rodrigues, A. S. L. (2004a). Species, data, and conservation planning, *Conservation Biology*, 18(6), 1682-1688.
- 9. Brooks, T. M., da Fonseca, G. A. B., Rodrigues, A. S. L. & (2004b). Protected areas and species, *Conservation Biology*, *18*(3), 616-618.
- Chen, G., Kéry, M., Plattner, M., Ma, K. & Gardner, B. (2013). Imperfect detection is the rule rather than the exception in plant distribution studies, *Journal of Ecology*, *101*(1), 183-191.
- Cushman, S. A. & Landguth, E. L. (2012). Multi-taxa population connectivity in the Northern Rocky Mountains, *Ecological Modelling*, 231, 101-112.

- Davies, Z. G. & Pullin, A. S. (2007). Are hedgerows effective corridors between fragments of woodland habitat? An evidence-based approach, *Landscape Ecology*, 22(3), 333-351.
- Delattre, T., Pichancourt, J. B., Burel, F. & Kindlmann, P. (2010). Grassy field margins as potential corridors for butterflies in agricultural landscapes: A simulation study, *Ecological Modelling*, 221(2), 370-377.
- Di Minin, E. & Moilanen, A. (2014). Improving the surrogacy effectiveness of charismatic megafauna with well- surveyed taxonomic groups and habitat types, *Journal* of *Applied Ecology*, 51(2), 281-288.
- Dolman, P. M. (2012). Mechanisms and processes underlying landscape structure effects on bird populations. In R. J. Fuller (Ed), *Birds and Habitat: Relationships in Changing Landscapes* (pp. 93-124), Cambridge University Press, Cambridge, UK.
- Dolman, P. M., Panter, C. J. & Mossman, H. L. (2012). The biodiversity audit approach challenges regional priorities and identifies a mismatch in conservation, *Journal of Applied Ecology*, 49(5), 986-997.
- Dorazio, R. M. & Royle, J. A. (2005). Estimating size and composition of biological communities by modeling the occurrence of species, *Journal of the American Statistical Association, 100*(470), 389-398.
- Eycott, A. E., Marzano, M. & Watts, K. (2011). Filling evidence gaps with expert opinion: The use of Delphi analysis in least-cost modelling of functional connectivity, *Landscape and Urban Planning*, 103(3-4), 400-409.
- Fens for the Future Partnership (2012) Fens for the Future a strategic plan for Fenland: a proposal for an enhanced ecological network, Fens for the Future Partnership, Peterborough,

UK: http://www.lincsfenlands.org.uk/index.php?page=BiodiversityFensFuture.

- Foster, G. N., Foster, A. P., Eyre, M. D. & Bilton, D. T. (1989). Classification of water beetle assemblages in arable fenland and ranking of sites in relation to conservation value, *Freshwater Biology*, 22(3), 343-354.
- Freckleton, R. (2002). On the misuse of residuals in ecology: regression of residuals versus multiple regression, *Journal of Animal Ecology*, 71, 42-545.

- Galpern, P., Manseau, M. & Fall, A. (2011). Patch-based graphs of landscape connectivity: A guide to construction, analysis and application for conservation, *Biological Conservation*, 144(1), 44-55.
- 23. Herzon, I. & Helenius, J. (2008). Agricultural drainage ditches, their biological importance and functioning, *Biological Conservation*, *141*(5), 1171-1183.
- Hill, J. K., Thomas, C. D., Fox, R., Telfer, M. G., Willis, S. G., Asher, J. & Huntley, B. (2002). Responses of butterflies to twentieth century climate warming: implications for future ranges, *Proceedings of the Royal Society B-Biological Sciences*, 269(1505), 2163-2171.
- 25. Hill, M. O., Preston, C. D. & Roy, D. B. (2004). PLANTATT-attributes of British and Irish plants: status, size, life history, geography and habitats, Centre for Ecology & Hydrology, Peterborough, UK.
- 26. Hill, M. O. (2011). Local frequency as a key to interpreting species occurrence data when recording effort is not known, *Methods in Ecology and Evolution*, *3*(1), 195-205.
- Isaac, N. J. B., van Strien, A. J., August, T. A., de Zeeuw, M. P. & Roy, D. B. (2014). Statistics for citizen science: extracting signals of change from noisy ecological data, *Methods in Ecology and Evolution*, 5(10), 1052-1060.
- Klar, N., Herrmann, M., Henning-Hahn, M., Pott-Doerfer, B., Hofer, H. & Kramer-Schadt, S. (2012). Between ecological theory and planning practice: (Re-) Connecting forest patches for the wildcat in Lower Saxony, Germany, *Landscape and Urban Planning*, *105*(4), 376-384.
- Krewenka, K. M., Holzschuh, A., Tscharntke, T. & Dormann, C. F. (2011). Landscape elements as potential barriers and corridors for bees, wasps and parasitoids, *Biological Conservation*, 144(6), 1816-1825.
- Kéry, M. (2011). Towards the modelling of true species distributions, *Journal of Biogeography*, 38(4), 617-618.
- 31. Lawson, C. R., Bennie, J. J., Thomas, C. D., Hodgson, J. A. & Wilson, R. J. (2012). Local and landscape management of an expanding range margin under climate change, *Journal of Applied Ecology*, 49(3), 552-561.

- 32. Lawson, C. R., Bennie, J. J., Thomas, C. D., Hodgson, J. A. & Wilson, R. J. (2014). Active management of protected areas enhances metapopulation expansion under climate change, *Conservation Letters*, 7(2), 111-118.
- Legendre, P. & Legendre, L. (2012). Numerical Ecology (3rd edition), In: *Developments in Environmental Modelling*, Elsevier, Amsterdam, The Netherlands.
- Leslie, A. W., Smith, R. F., Ruppert, D. E., Bejleri, K., McGrath, J. M., Needelman, B. A. & Lamp, W. O. (2012). Environmental factors structuring benthic macroinvertebrate communities of agricultural ditches in Maryland, *Environmental Entomology*, 41(4), 802-812.
- Liira, J. & Paal, T. (2013). Do forest-dwelling plant species disperse along landscape corridors? *Plant Ecology*, 214(3), 455-470.
- Lombard, A. T., Cowling, R. M., Pressey, R. L. & Rebelo, A. G. (2003). Effectiveness of land classes as surrogates for species in conservation planning for the Cape Floristic Region, *Biological Conservation*, *112*(1-2), 45-62.
- MacKenzie, D. I. & Kendall, W. L. (2002). How should detection probability be incorporated into estimates of relative abundance? *Ecology*, *83*(9), 2387-2393.
- Milsom, T. P., Sherwood, A. J., Rose, S. C., Town, S. J. & Runham, S. R. (2004). Dynamics and management of plant communities in ditches bordering arable fenland in eastern England, *Agriculture Ecosystems & Environment*, 103(1), 85-99.
- Morton, R. D., Rowland, C., Wood, C., Meek, L., Marston, C., Smith, G., Wadsworth, R. & Simpson, I. (2011). Land Cover Map 2007 (1km raster dominant Target Class, GB), NERC-Environmental Information Data Centre, Swindon, UK.
- 40. Mossman, H. L., Panter, C. J. & Dolman, P. M. (2012). Fens Biodiversity Audit, Report to Fens for the Future steering group, Peterborough, UK.
- Mountford, O. & Arnold, H. (2006). Aquatic plant diversity in arable ditches: scoping study, Centre for Ecology and Hydrology, Monks Wood, UK.
- 42. Natural England (2013). National Character Area profile: 46. The Fens, Natural England, Peterborough, UK, http://www.naturalengland.org.uk/publications/nca/the_fens.aspx.
- Noss, R. F. (1990). Indicators for monitoring biodiversity a hierarchical approach, *Conservation Biology*, 4(4), 355-364.

- Nowicki, P., Vrabec, V., Binzenhoefer, B., Feil, J., Zaksek, B., Hovestadt, T. & Settele, J. (2014). Butterfly dispersal in inhospitable matrix: rare, risky, but long-distance, *Landscape Ecology*, 29(3), 401-412.
- Ockenden, M. C., Deasy, C., Quinton, J. N., Bailey, A. P., Surridge, B. & Stoate, C. (2012). Evaluation of field wetlands for mitigation of diffuse pollution from agriculture: sediment retention, cost and effectiveness, *Environmental Science & Policy*, 24, 110-119.
- 46. Palmer, M., Drake, M. & Stewart, N. (2013). A manual for the survey and evaluation of the aquatic plant and invertebrate assemblages of grazing marsh ditch systems. Buglife, Peterborough, UK.
- 47. R Core Team (2012). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. ISBN 3-900051-07-0, URL http://www.R-project.org/.
- 48. Rayfield, B., Fortin, M.-J. & Fall, A. (2011). Connectivity for conservation: a framework to classify network measures, *Ecology*, *92*(4), 847-858.
- Saura, S., Bodin, O. & Fortin, M.-J. (2014). Stepping stones are crucial for species' longdistance dispersal and range expansion through habitat networks, *Journal of Applied Ecology*, *51*(1), 171-182.
- 50. Schindler, S., von Wehrden, H., Poirazidis, K., Wrbka, T. & Kati, V. (2013). Multiscale performance of landscape metrics as indicators of species richness of plants, insects and vertebrates, *Ecological Indicators*, *31*, 41-48.
- Simon, T. N. & Travis, J. (2011). The contribution of man-made ditches to the regional stream biodiversity of the new river watershed in the Florida panhandle, *Hydrobiologia*, 661(1), 163-177.
- 52. Twisk, W., Noordervliet, M. A. W. & Ter Keurs, W. J. (2000). Effects of ditch management on caddisfly, dragonfly and amphibian larvae in intensively farmed peat areas, *Aquatic Ecology*, 34(4), 397-411.
- 53. Virkkala, R., Luoto, M., Heikkinen, R. K. & Leikola, N. (2005). Distribution patterns of boreal marshland birds: modelling the relationships to land cover and climate, *Journal of Biogeography*, 32(11), 1957-1970.

- 54. Williams, P., Whitfield, M., Biggs, J., Bray, S., Fox, G., Nicolet, P. & Sear, D. (2004). Comparative biodiversity of rivers, streams, ditches and ponds in an agricultural landscape in Southern England, *Biological Conservation*, *115*(2), 329-341.
- 55. Zulka, K. P., Abensperg-Traun, M., Milasowszky, N., Bieringer, G., Gereben-Krenn, B.-A., Holzinger, W., Hoelzler, G., Rabitsch, W., Reischuetz, A., Querner, P., Sauberer, N., Schmitzberger, I., Willner, W., Wrbka, T. & Zechmeister, H. (2014). Species richness in dry grassland patches of eastern Austria: A multi-taxon study on the role of local, landscape and habitat quality variables, *Agriculture Ecosystems & Environment, 182*, 25-36.

TABLE LEGENDS

Table 1. Definition and data source of environmental predictors used to model the distribution ofFens biodiversity.

Table 2. Variation in the richness of wetland groups of species explained by the minimum models.

Table 1. Definition and data source of environmental predictors used to model the distribution of Fens biodiversity.

	Abbreviation used in	Source	
Environmental predictor	Figure 2		
Mean elevation above sea level: mean elevation of all	Elevation		
50 m x 50 m cells within the 1 km square			
Distance to nearest fenland island: Fenland island	Distance to Fen Island	- Edina Digimap	
defined as areas >0.1 km ² with an elevation of \ge 5 m		Ordinance Survey	
(excluding coastal cliffs at Skegness and islands within		(OS) PANORAMA	
large urban areas). Several large 'islands' within 1000 m		DTM (Digital Terrain	
of the fenland basin were incorporated into the basin,			
i.e. not considered islands.		Model) 1:50,000, 50m _ cells	
Distance to fenland basin: basin was defined as the 5 m			
contour boundary, unless the area had been defined as a			
fenland island.			
Presence of either an A or B road within a square	Presence of a road	OS Meridian 2 (1:50	
		000)	
Distance to nearest SSSI comprising wetland habitats	Distance to wetland	Natural England GIS	
	SSSI	Digital Boundary	
		Datasets	
Percentage of square comprising urban areas. Urban	% urban		
defined from OS Strategic 1:250,000		- Edina Digimap	
Network distance along 'ditches' to the nearest 'main	Distance to river	Ordinance Survey	
river'/coastline: calculated using network cost distance.		Strategic 1:250,000	
Ditch was defined using the VectorMap District		Strategie 1.250,000	
Surface_Water polyline for accurate mapping of small		VectorMap District	
ditches and open water, and the Tidal_Boundary		(1:25,000)	
(High/Low Water Mark) polyline because the surface		(1.23,000)	
water data stop at the tidal boundary.			

Network distance along ditch/river to the tidal	Distance to tidal	-
boundary: calculated using network cost distance (see	boundary	
below for full description). Ditch/river defined using the		
Edina Digimap River_polyline and VectorMap District		
Surface_Water polyline. Tidal boundary was defined as		
the high water mark using the VectorMap District		
<i>Tidal_Boundary</i> polyline.		
Index length of all ditches per 1 km square: ditches were	Length of surface	-
defined as above. This is considered an index because	water	
polylines defined each bank of wide ditches or rivers,		
resulting in double-counting, as such the lengths are not		
accurate.		
Percentage of rough and neutral grassland	% rough/neutral	Land Cover Map
	grassland	2007. Centre for
		Ecology and
		Hydrology
Percentage of grades of Agricultural Land	% grades 3 & 4	Natural England GIS
Classification: summed percentage area of grades 3 and		Digital Boundary
4		Datasets
Percentage of each peat soil type defined using	% peat; % peaty	
Cranfield Soil Class; Peat; Seasonally wet deep peat to	loam; % deep sand	
loam; Seasonally wet deep clay over peat (marine	over peat; % deep clay	
alluvium and fen peat) and Seasonally wet deep sand	over peat	NATMAP Cranfield
over peat (glaciofluvial drift and peat).		University
Percentage of silt soil, defined as the Cranfield Soil	% silt	
Class "Seasonally wet deep silty"		
Occurrence of a county flora: 0/1 if in a flora recorded	County flora	
county		

		% variation of total r ² explained		
	Total r ²		Independent	Joint effect of
		Independent effect	effect of	recording and
		of recording effort	environment	environment
Odonata	30.4	27.3	52.8	20.0
Fen Specialists	27.2	46.4	27.1	26.4
Aquatic species	30.5	49.2	35.5	15.3
Aquatic plants	59.8	75.4	19.9	4.8
Littoral species	31.5	64.4	33.1	2.5
Wetland species	39.0	57.8	25.5	16.7
Wetland plants	63.9	76.2	17.1	6.7

Table 2. Variation in species richness explained by the minimum models.

LIST OF FIGURES

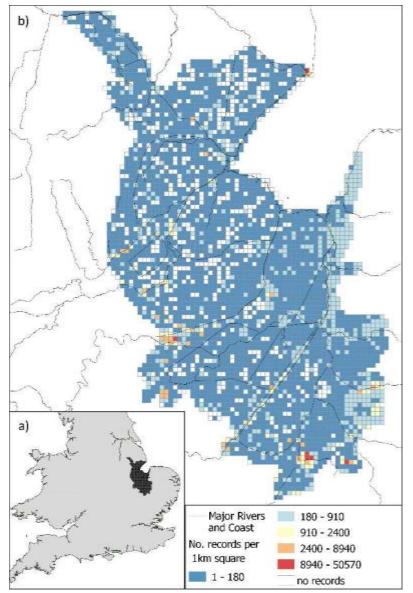
Figure 1. (a) The location of the Fens region within the UK, and b) the intensity of recording effort within the Fens, shown as number of records per 1-km square. Class intervals calculated using jenks natural breaks.

Figure 2. Mean (\pm SE) standardised effect size (β values) of environmental predictor variables on the richness of ditch indicator groups. Only significant (p<0.05) effects are shown. Predictor abbreviations are provided in Table 1.

Figure 3. Predicted richness per 1-km square for a) Odonata species, b) littoral priority species, c) aquatic priority species, d) aquatic plants, e) wetland priority species, f) wetland plants, and g) Fens Specialists. White areas denote 1 km squares that were excluded from models. Class intervals calculated using jenks natural breaks.

Figure 4a. Mean of the ranks of predicted species richness per 1 km square of the seven wetland biological indicator groups. A low rank (high number) is given to squares with low predicted biodiversity and high rank (tied, highest = 44) to areas with high biodiversity. Main rivers and (a) connectivity corridors proposed by the Fens for the Future Partnership (excluding the Landscape Corridor) (FFFP 2012), and (b) connectivity corridors re-routed through areas of higher wetland species richness are shown. White areas denote 1 km squares excluded from models. Class intervals calculated using jenks natural breaks.

Figure 5. Length (km) of primary and secondary connectivity corridors originally proposed by the Fens for the Future Partnership and alternative corridors selectively routed through areas of predicted higher wetland richness. Bars are shaded according to quartiles of the mean of ranks of biodiversity richness per 1-km square across seven indicator groups (Q1: 44-950, Q2: 951-1561, Q3: 1562-2372, Q4: 2373-3000).



Contains Ordnance Survey data © Crown copyright and Database Right 2014.

Figure 1. (a) The location of the Fens region within the UK, and b) the intensity of recording effort within the Fens, shown as number of records per 1-km square. Class intervals calculated using jenks natural breaks.

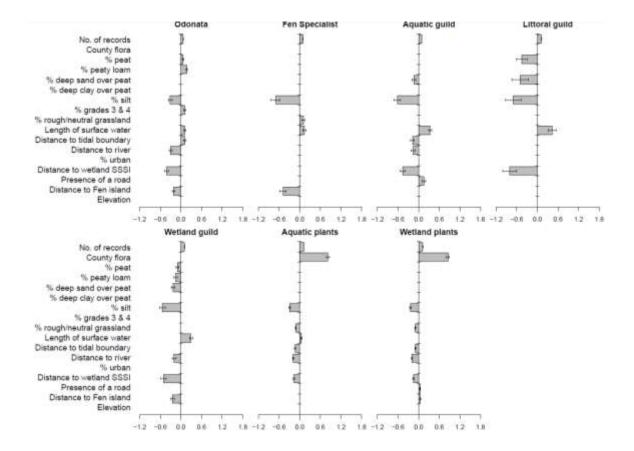


Figure 2. Mean (\pm SE) standardised effect size (β values) of environmental predictor variables on the richness of ditch indicator groups. Only significant (p<0.05) effects are shown. Predictor abbreviations are provided in Table 1.

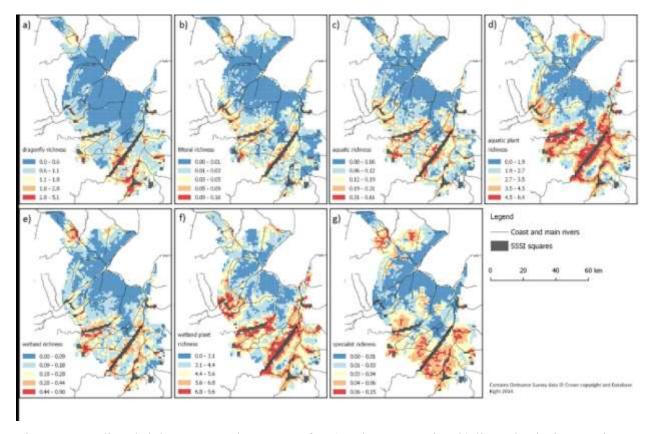
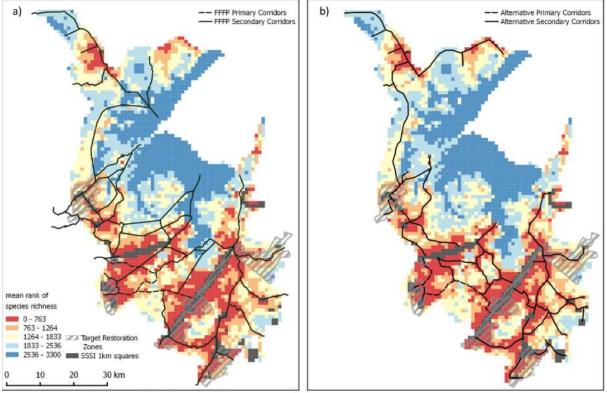


Figure 3. Predicted richness per 1-km square for a) Odonata species, b) littoral priority species, c) aquatic priority species, d) aquatic plants, e) wetland priority species, f) wetland plants, and g) Fens Specialists. White areas denote 1 km squares that were excluded from models. Class intervals calculated using jenks natural breaks.



Contains Ordnance Survey data © Crown copyright and Database Right 2014.

Figure 4a. Mean of the ranks of predicted species richness per 1 km square of the seven wetland biological indicator groups. A low rank (high number) is given to squares with low predicted biodiversity and high rank (tied, highest = 44) to areas with high biodiversity. Main rivers and (a) connectivity corridors proposed by the Fens for the Future Partnership (excluding the Landscape Corridor) (FFFP 2012), and (b) connectivity corridors re-routed through areas of higher wetland species richness are shown. White areas denote 1 km squares excluded from models. Class intervals calculated using jenks natural breaks.

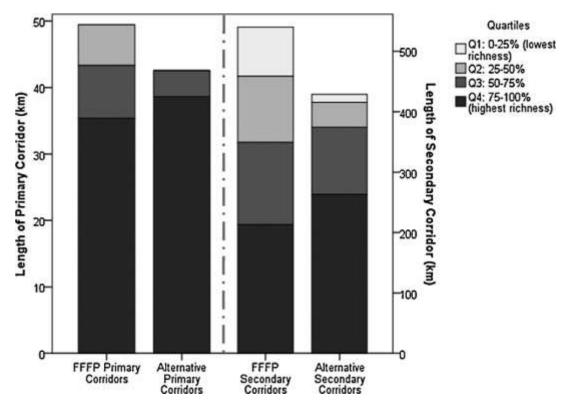


Figure 5. Length (km) of primary and secondary connectivity corridors originally proposed by the Fens for the Future Partnership and alternative corridors selectively routed through areas of predicted higher wetland richness. Bars are shaded according to quartiles of the mean of ranks of biodiversity richness per 1-km square across seven indicator groups (Q1: 44-950, Q2: 951-1561, Q3: 1562-2372, Q4: 2373-3000).