



Vegetation diversity of conventional and organic hedgerows in Denmark

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Received 26 November 2002; received in revised form 21 March 2003; accepted 26 March 2003

Abstract

Many attempts have been made to reduce the impact of modern conventional farming on the environment and semi-natural ecosystems. One of them is organic farming, known primarily for the absence of pesticides and artificial fertilisers. The objective of this study was to study and test the differences in the spontaneous vegetation of comparable hedgerows in the same area situated within organic and conventional farming systems. The hedge bottom vegetation was surveyed during August 2001 in 13 hedgerows of each farming system. Farming type had not changed on either side of the hedgerows for the lifetime of the hedges (10–14 years). Sampling was associated with a set of 16 measured environmental variables. In the two farming systems hedgerows were comparable in terms of landscape, age, soil type, nutrient status and width. A mixed analysis of variance (ANOVA) found no significant difference in measured soil and radiation variables between farming types. Farming types only differed in the use of pesticides. Significant differences between farming types in plant species diversity at alpha, beta and gamma levels were found. More species that are normal in semi-natural habitats were found on organic farms. There was an overlap in species composition between farming type, but a slightly higher species turnover on conventional farms. The ordination axes were highly correlated with calibrated Ellenberg values of fertility, light and soil moisture. Soil fertility and farming type were important factors to explain variation in species composition. Organic farming had a significantly reduced impact on hedge bottom vegetation compared to conventional farming. Higher extinction due to pesticide drift and immigration rates may be responsible for the significantly higher species diversity and different species composition in hedges on organic farms. The differences in species diversity and plant types are briefly discussed.

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Keywords: Farming type; Fertility; Hedge bottom; Naturalness; Pesticide; Species diversity

1. Introduction

Hedges have a multitude of functions (Baudry et al., 2000) and are receiving increasing attention as refuges

for wildlife in countries with intensive agricultural production. Most studies on hedgerows originate from the UK, France and Canada (McCollin et al., 2000; Barr and Petit, 2001; Boutin et al., 2001). The ecological function of hedgerows in the agricultural landscape matrix has received increased attention since the 1980s (Baudry et al., 2000; Le Coeur et al.,

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2002) and spontaneous vegetation in hedgerows can be considered as part of a metapopulation matrix. The function of hedgerows as dispersal corridors of forest species (McCollin et al., 2000; Smart et al., 2001) and refuges of grassland species (Smart et al., in press) has recently been described. Likewise, the neighbourhood of potential sources for species immigrating to hedges has also been investigated (Mercer et al., 1999; Boots, 2001).

Neighbouring land-use is known to exert a strong impact on hedgerow vegetation, especially on the hedge bottom flora (Boatman et al., 1994; Cummins and French, 1994; Hegarty et al., 1994; McAdam et al., 1994; Le Coeur et al., 2002) and other linear biotopes (De Snoo and van der Poll, 1999; Hald, 2002). Two of the major impact factors of modern conventional agriculture on existing habitats are displacement of fertilisers and pesticides drift (Pollard et al., 1974; Timmermans and Cuppen, 1988; Boatman, 1992; Hegarty et al., 1994; Baudry et al., 2000; McCollin et al., 2000). Fertilisation is held as the most influential factor (Hegarty et al., 1994) and its impact has been illustrated for a range of semi-natural vegetation types (Traczyk and Kotowska, 1976; Virtanen et al., 2000) including hedge bottom vegetation (Tsiouris and Marshall, 1998). The boundary vegetation in intensive agricultural areas is characterised by a relatively species-poor, nitrophilous vegetation (Hegarty et al., 1994; Kleijn and Verbeek, 2000; Tybirk et al., 2001). The drift of pesticides has also been found to influence the organisms of hedges (Hald et al., 1994; Jobin et al., 1997; Boutin and Jobin, 1998), and its impact on non-crop species is well documented (Marrs et al., 1989; Marrs and Frost, 1997). The influence of pesticide drift is also well known for other field boundary vegetation (Fischer and Milberg, 1997; Kleijn and Snoeiijing, 1997; De Snoo, 1999) and semi-natural vegetation (Marrs et al., 1989). The use of fertiliser (Keeney and Hatfield, 2001) and pesticides (Boutin and Rogers, 2000) has increased tremendously since the Second World War and it is not clear whether fertilisation and pesticide drift interact or whether pesticides alone affect the hedge bottom vegetation.

Continuity in time and space is very important in terms of species composition and botanical diversity in hedges (Warming, 1919; Pollard et al., 1974; Dowdeswell, 1987; Bunce et al., 1994; Tybirk et al.,

2001). Some studies indicate that the age effect on tree diversity is quite weak (Willmot, 1980). Nevertheless, the removal of older hedgerows is an irreversible process since new hedgerows contain different vegetation compared to old ones (Boutin et al., 2001). In Great Britain, management plays an important role for hedgerow vegetation (Moonen and Marshall, 2001).

To reduce the impact of modern conventional farming on semi-natural ecosystems buffer zones, such as conservation headlands, unsprayed field margins (Marshall and Moonen, 2002) and selective plant control (Boatman, 1989) have been shown to improve or at least preserve the natural biodiversity (Schumacher, 1984; Hald et al., 1994; De Snoo, 1997; De Snoo and de Wit, 1998). Integrated (Ogilvy et al., 1995) and organic farming system (Lampkin, 1990) are also ways to reduce the impact on semi-natural ecosystems. Organic farming covers today about 3% of the European agricultural land (Yussefi and Willer, 2003). While there are indications of a positive influence on biodiversity conservation on cultivated fields and uncultivated semi-natural biotopes (Stolze et al., 2000; Azzez, 2000; Mäder et al., 2002), the impact on semi-natural biotopes is poorly documented. Likewise there is hardly any evidence of long-term effects of organic farming on semi-natural species assemblages in hedgerows (Tybirk and Ejrnæs, 2001). This study aimed at investigating the potential influence of conventional and organic farming on adjacent hedgerow vegetation during the first 10–14 years after the hedgerow was planted and to test the following null hypotheses: (1) the type of farming (organic versus conventional) does not affect alpha, beta and gamma plant species diversity and (2) it does not influence species composition or plant types in hedge bottom vegetation.

2. Methods and material

Thirteen organic hedges on five organic farms and 13 conventional hedges on eight conventional farms were investigated. Organic hedgerows were selected first and then in the conventional hedgerows were randomly selected in the same district and, where possible on the same roads, at a median distance of 850 m. All hedges were located in the southwestern part of Jutland on sandy Saalian moraines or Weichelian outwash

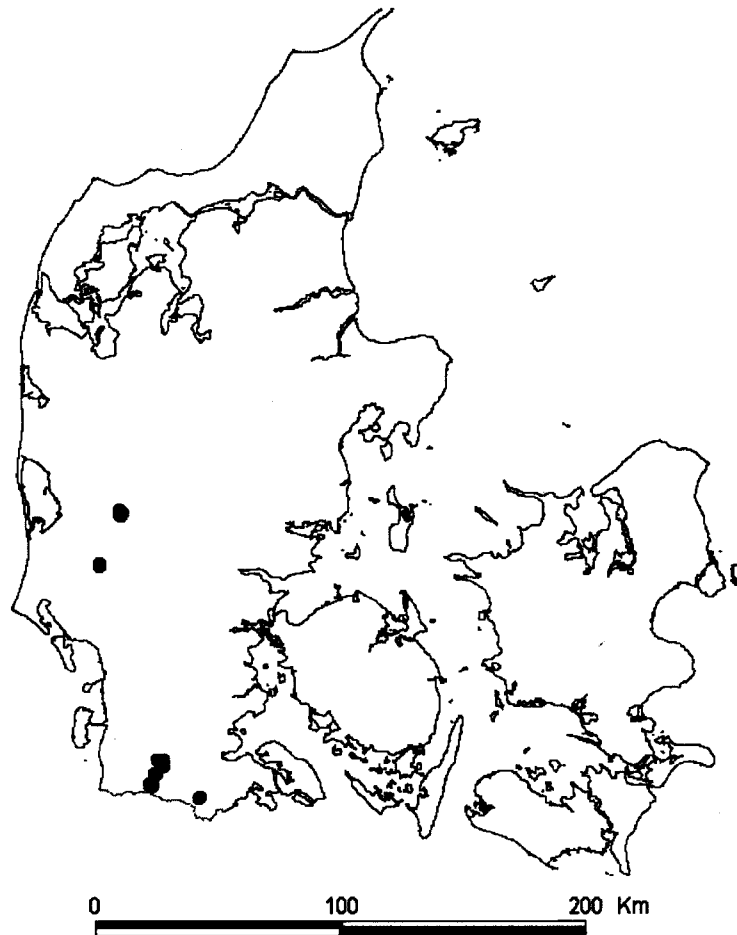


Fig. 1. Map of Denmark, the dots showing location of sampling sites.

136 plains (Fig. 1). Average temperature for January and
 137 July was -0.1 and 15.2 °C, respectively and yearly av-
 138 erage precipitation 823 mm (Frich et al., 1997; Laursen
 139 et al., 1999).

140 Hedgerows included on organic farms had been es-
 141 tablished and cultivated without any pesticide. Neigh-
 142 bouring fields on both sides had also been managed
 143 organically or conventionally since establishment of
 144 the hedgerow. Hedgerows had agricultural fields on
 145 both sides and none of the farmers used headlands or
 146 unsprayed field margins. All hedgerows had been es-
 147 tablished 10–14 years ago by a company (Hedeselska-
 148 bet), using the same technique and all contained three
 149 to four rows of a mixture of exotic and native decid-
 150 uous woody species. The planting distance between
 151 mixtures of 15 and 25 woody species was 1×1.2 m

152 with a minimum hedge length of 120 m and plough-
 153 ing to a depth of 0.5 m to avoid seed from the seed
 154 bank. The starting point was hence similar and the
 155 recorded species have colonised from the surrounding
 156 landscape.

157 Data were collected in August 2001 of six sampling
 158 plots (three pairs) of 10 square metre ($0.5 \text{ m} \times 20 \text{ m}$)
 159 placed along each hedgerow, 20 m apart. In each pair
 160 one plot was placed 0.2 m from the field edge, and one
 161 was placed in the centre of the hedgerow, between the
 162 first and second planted row. All vascular plants and
 163 bryophytes were recorded in each 10 square metre
 164 plot and species abundance was evaluated by the num-
 165 ber of rooted species (soil contact for bryophytes) in
 166 10 Raunkjær circles of $1/10$ square metre (Raunkjær,
 167 1910). Other species outside the Raunkjær circles

168 but inside the plot were noted and given the lowest
169 frequency score (=1). Planted woody species and
170 their regeneration were not included. A total list of
171 species from 120 m on one side of the hedgerows was
172 obtained by noting supplementary species between
173 plots. To optimise comparison between hedges the
174 most south-orientated was considered.

175 Photosynthetic active radiation (PAR) was mea-
176 sured with a LI-191SA Line Quantum Sensor
177 (LI-COR 1991). The sensor was 1 m long, and double
178 measurements were performed at a height of 0.5 m
179 in three positions along the centre of each plot. Mea-
180 surements were expressed as a percentage of open
181 field measurements perpendicular to the sample plot.

182 The orientation of the hedgerows was measured
183 with a compass. Hedge bank height was measured
184 in two positions along the sample plot and aver-
185 aged. Hedge width was measured at both ends of the
186 hedgerows. Height of hedges was estimated in four
187 classes (<3 m, 3–5 m, 5–7 m and >7 m).

188 Soil samples from each sampling plot were col-
189 lected in October 2001. Five sub-samples of soil
190 were collected 0–5 cm below the A₀-horizon, lumped
191 and sifted through a 2 mm sieve to remove roots and
192 stones. Water content of the soil was measured gravi-
193 metrically. Soil pH and conductivity were measured
194 in mixtures of soil and H₂O (1:5 v/v). Fresh soil
195 (10 g) was immediately extracted for 1 h in 50 ml of
196 0.5 mol/l K₂SO₄ to recover soil inorganic N and P.
197 The extracts were filtered through Whatman GF-D
198 filters and frozen until their NH₄⁺-N content could
199 be analysed with the indophenol method, NO₃⁻-N
200 with the cadmium reduction method (Allen, 1989).
201 A fraction of each soil sample was dried at 70 °C
202 to constant weight and finely ground. About 4 g of
203 the dried soil was burnt at 550 °C for determina-
204 tion of the organic content. Another fraction, about
205 200 mg dry soil, was digested in a sulphuric acid
206 and selenous acid mixture for 1 h (Kedrowski, 1983) to
207 analyse total P with the molybdenum blue method
208 and total N with the indophenol method, using a Hi-
209 tachi U-2000 spectrophotometer. Soil carbon content
210 (total C) was measured with a LECO CNS-1000 ac-
211 cording to Sørensen and Bülow-Olsen (1994). Soil
212 texture was finally obtained from the marginal plots
213 and centre plots by lumping the soils from the three
214 central plots and three marginal plots resulting in two
215 samples per hedgerow. Soil was sieved to find the

proportion of coarse sand (200–2000 µm), fine sand 216
(63–200 µm), coarse silt (20–63 µm), silt (2–20 µm) 217
and clay (<2 µm) (Sørensen and Bülow-Olsen, 1994). 218
Coarse sand and fine sand were correlated and there- 219
fore only coarse sand was used. Every vegetation 220
sample was associated with a set of 16 environmental 221
variables (Table 1). 222

The identification and taxonomy of vascular plants 223
was based on Hansen (1993), Hubbard (1984), 224
Pedersen and Schou (1988) and for mosses on 225
Andersen et al. (1976). 226

2.1. Data analysis 227

Plant species diversity in the hedgerow was calcu- 228
lated on three levels: α (plot level), β (species turnover) 229
and γ (hedge level). A mixed model analysis of vari- 230
ance (ANOVA) was used to test differences in al- 231
pha and gamma diversity between management types. 232
ANOVA was also used to test differences in explana- 233
tory variables and plant types between management 234
types. In both cases random variables were hedge and 235
block number within hedge. Fixed variables were man- 236
agement type and position in hedgerow (central or 237
boundary plot). The overall statistical model applied 238
was 239

$$Y_{ijkl} = \mu + \alpha_i + \beta_j + (\alpha\beta)_{ij} + C_{ik} + D_{ikl} + \varepsilon_{ijkl} \quad 240$$

where μ is the mean, α_i denotes the effect of manage- 241
ment system i , β_j is the effect of position in hedgerow 242
(j , boundary or central), $(\alpha\beta)_{ij}$ represents the interac- 243
tion between management and position, C_{ik} is the ef- 244
fect of hedgerow number k in management system i , 245
 D_{ikl} represents the effect of pair (block) l in hedgerow 246
 k in management system i , and ε_{ijkl} is the residual ef- 247
fect. C_{ik} was assumed to be mutually independent and 248
normally distributed with a mean of 0 and a variance 249
of σ_B^2 , ε_{ijkl} being assumed to be mutually independent 250
and normally distributed with a mean of 0 and a vari- 251
ance of σ^2 , D_{ikl} is mutually independent and normally 252
distributed with a mean of 0 and a variance of σ_D^2 . 253

The use of beta diversity followed Whittaker (1972) 254
and Ejrnæs et al. (2002). For comparison of the two 255
farming types, 12 random subsets of 10 sample plots 256
were drawn from each farming type, and beta diversity 257
was calculated as the total number of species in the 10 258
sample plots divided by the average number. The t -test 259
was used to compare beta diversity per farming type. 260

Table 1

Explanatory variables and variance explained ranked according to percent of explained variation (sixth column) evaluated by running forward selection of all variables followed by Monte Carlo Permutation test

Explanatory variables	Units	Range (minimum and maximum)	Mean	Trans-formations	Variance explained by this variable	Level of significance
Total P	mg/g	0.036–0.857	0.283	\sqrt{x}	12.0	***
Location	Nominal	0–1	–	–	10.4	***
Total C	g/100 g soil	0.842–9.386	3.22	$\log(x)$	10.2	***
Coarse silt	g/100 g soil	1–9.1	2.09	$\log(x)$	9.5	***
Type of farming (organic vs. conventional)	Nominal	0–1	–	–	9.4	***
Bank height	m	0–0.8	0.086	\sqrt{x}	7.5	***
Coarse sand	g/100 g soil	40.1–81.1	66.88	–	7.0	***
Clay	g/100 g soil	2.5–8.9	3.96	$\log(x)$	5.5	***
Total N	mg/g	0.072–3.729	1.493	\sqrt{x}	4.6	**
Conductivity	MicroSiemens/cm	14.6–74.8	32.88	$\log(x)$	4.3	**
Silt	g/100 g soil	0.9–6.8	2.87	$\log(x)$	3.9	*
Loss on ignition	%	1.5–9.8	6.31	–	3.7	**
Orientation	°	60–260	137	$\log(x)$	3.5	*
pH _{H₂O}	[H ⁺]	4.3–6.85	5.43	$\exp(x)$	3.5	*
Light (average)	Percentage of field	0.5–78.5	15.23	$\log(x)$	3.2	NS
Water content	Percentage of dry weight	5.4–35.5	17.0	$\log(x)$	1.7	NS

NS: not significant.

* *P* values between 0.01 and 0.05.

** *P* values between 0.001 and 0.01.

*** *P* values <0.001.

261 The vegetation data comprised 156 plots with 144
 262 taxa. Prior to multivariate analysis 71 species with
 263 less than four occurrences were deleted according to
 264 Økland (1990). A detrended correspondence analysis
 265 (DCA) (Hill, 1979) was performed with default op-
 266 tions, that is, detrending by 26 segments, non-linear
 267 rescaling and rescaling of axes for species abundance
 268 data. All explanatory variables were examined for kur-
 269 tosis and skewness. Variables with numerical values
 270 >1 were transformed to reduce kurtosis and skewness.
 271 The type of transformations that resulted in the small-
 272 est kurtosis and skewness value was chosen (Table 1).
 273 Hereafter variables were centred and standardised
 274 to obtain unit variance. A constrained ordination
 275 (CCA) was used to estimate the explanatory value
 276 of the variables followed by forward selection and
 277 Monte Carlo Permutation test with 999 unrestricted
 278 permutations (Ter Braak and Smilauer, 1997–1999).
 279 The indicator values of Ellenberg et al. (1991) for
 280 temperature (*T*), moisture (*F*), fertility (*N*), light (*L*)
 281 and acidity (*R*) were used to calculate a weighted
 282 mean for all plots. Product–moment correlations were

calculated between ordination axes and explanatory 283
 variables. 284

3. Results 285

286 Out of 26 hedgerows, 12 were adjacent to ley (5 286
 287 conventional and 7 organic) with a typical lifetime of 287
 288 2–3 years, and 14 adjacent to barley, rape and veg- 288
 289 etables (8 conventional and 6 organic). Mean hedge 289
 290 width (furrow to furrow) was 5.2 m (S.E. = 0.19), and 290
 291 the median height class was 3 (5–7 m). All hedgerows 291
 292 had been established in a standardised way in the 292
 293 same period and have a very similar and comparable 293
 294 structure. Thus, width and height data were not 294
 295 included in the following analysis, but are mentioned 295
 296 for comparison with other countries (Baudry et al., 296
 297 2000). The mean orientation of organic and conven- 297
 298 tional hedgerows was 148° (SSE) and 125°(ESE), the 298
 299 mean bank heights were 0.06 and 0.12 m, respectively 299
 300 and were the only factors for which mean values dif- 300
 301 fered significantly between farming types (Table 2). 301

Table 2

Mean values of explanatory variables and results of mixed analysis of variance of transformed values

Explanatory variables	Mean values		Component of variance (random effects)			P-values		
	Organic	Conventional	Hedge number	Block	Residual	Type of farming	Position	Type of farming × position
pH _{H2O}	5.40	5.46	0.4	0.1	0.5	0.6	<0.0001	0.7
Conductivity	34.25	31.52	0.5	0.004	0.5	0.4	0.1	0.07
Water content	23.11	18.17	0.6	0.04	0.4	0.08	0.9	0.3
Total P	0.30	0.27	0.8	0	0.2	0.6	<0.0001	0.4
Total N	1.55	1.44	0.5	0	0.5	0.7	0.005	0.9
Loss on ignition	4.9	5.3	0.7	0.08	0.2	0.2	0.1	0.8
Coarse sand	65	69	0.9	–	0.09	0.3	0.002	0.03
Coarse silt	2.1	2.0	0.9	–	0.2	0.9	<0.0001	0.03
Clay	4.2	3.8	1.0	–	0.09	0.5	0.007	0.2
Silt	3.2	2.6	0.7	–	0.2	0.2	<0.0001	0.2
Total C	3.7	2.8	0.9	–	0.08	0.2	<0.0001	0.3
Light	14	17	0.3	0	0.3	0.2	<0.0001	0.3
Bank height	0.06	0.12	6.0	0.3	0	<0.0001	1.0	1.0
Orientation	148	125	0.3	0	0	<0.0001	1.0	1.0

Estimated relative importance of random variables and level of significance are shown. "Position" refers to inside and outside the hedgerow.

302 Median values were equal for both bank height (zero)
303 and orientation (120°).

304 A total of 144 plant species were found of which 14
305 were spontaneous woody, 118 herb and 12 bryophyte
306 species, 101 of which occurred in conventional, 128
307 in organic hedgerows. Ecological classification and
308 preference of species were evaluated according to
309 Hansen (1993) and Smith (1978). Of all the species,
310 73 were found in semi-natural habitats, 40 being ex-
311 clusively found in semi-natural habitats, 12 of which
312 being forest species. Seventy-one species were con-

sidered as arable and ruderal species, one protected 313
species was found (*Epipactis helleborine* (L.) Crantz.) 314
in an organic hedgerow. Total numbers of hedge bot- 315
tom species (gamma diversity) ranged from 24 to 53 316
(\bar{x} = 38.7, S.E. = 1.6) in organic, from 20 to 36 317
(\bar{x} = 28.8, S.E. = 1.1) in conventional hedgerows, 318
the difference being highly significant (Table 3). The 319
difference in average species richness of sample plots 320
(alpha diversity) was also highly significant (organic 321
15.1 versus conventional 12.5, P = 0.013), boundary 322
samples also being significantly different (organic

Table 3

Analysis of variance of plant types and species diversity between organic and conventional farming (estimated relative importance of random variables made on presence-absence data in all sampling plots, with P -values for fixed variables)

	Mean values in plots		Component of variance (random effects)			P-values		
	Organic	Conventional	Hedge number	Block	Residual	Type of farming	Position	Type of farming × position
Weed and ruderal species	8.3	6.4	4.9	0	6.2	0.041	<0.0001	0.0056
Semi-natural species	5.1	3.9	1.4	0	2.0	0.018	0.0041	0.236
Forest species	0.41	0.47	0.1	0	0.3	0.82	<0.0001	0.79
Bryophytes	0.96	0.65	0.3	0.03	0.4	0.33	<0.0001	0.79
Alpha diversity total	15.1	12.5	7.3	0	11.9	0.013	0.0003	0.006
Alpha diversity (outside)	17.2	12.8	12.8	6.5	1.0	0.0064	–	–
Alpha diversity (central)	13.7	12.2	12.5	5.1	1.0	0.33	–	–
Gamma diversity	38.7	28.8	–	–	–	0.0011	–	–

"Position" refers to inside and outside the hedgerow.

Table 4

Mixed analysis of variance of weighted mean values of Ellenberg et al. (1991) values for all sampling plots (estimated relative importance of random variables and *P*-values of fixed variables is given)

	Component of variance (random effects)			<i>P</i> -values		
	Hedge number	Block	Residual	Type of farming	Position	Type of farming × position
Light	0.03	0	0.04	0.87	<0.0001	0.12
Temperature	0.02	0	0.02	0.34	<0.0001	0.1585
Soil moisture	0.02	0	0.08	0.53	<0.0001	0.048
pH	0.7	0	1.3	0.09	0.38	0.95
Nitrogen	0.3	0.02	0.1	0.52	<0.0001	0.31

323 17.2 versus conventional 12.8, $P = 0.0064$). There
 324 was no difference between species richness in samples
 325 from inside the hedgerow (organic 13.7 versus con-
 326 ventional 12.2, $P = 0.33$). Organic hedgerows were
 327 significantly richer in weedy, ruderal and semi-natural
 328 species but there was no difference in frequency of
 329 bryophytes and forest species (Table 3). There was
 330 no significant difference in soil variables between the
 331 farming types (Table 2), whereas samples taken at the
 332 field edge and centrally in the hedgerows differed for
 333 most soil parameters.

334 There was no significant difference in weighted
 335 mean Ellenberg values between farming types
 336 (Table 4). The overall correlation between species
 337 richness of sampling plots and explanatory variables
 338 was highest for soil variables and insignificant for
 339 variables concerning hedge structure (Table 5). There
 340 was a significantly higher beta diversity (Whittaker,
 341 1972) on conventional farms (t -test, d.f. = 11, $P =$
 342 0.0074). The contribution of block number within the
 343 hedge to the total random variance was low compared
 344 to hedge number and residual effect (Tables 2–4)
 345 which means that differences within hedges were
 346 small compared to differences between hedges.

347 Species occurring in less than four samples were
 348 deleted from the data matrix, leaving 156 samples with
 349 73 species. Gradient lengths were 5.1, 5.2 and 3.4 for
 350 DCA axes 1–3. It is obvious from Fig. 2 that there was
 351 a great overlap in species composition between the
 352 two management types. DCA1 sample scores had the
 353 highest negative correlation with soil nutrient status
 354 (Ellenberg *N* value, *P* total and amount of coarse sand)
 355 and measured light variables (Table 5), indicative of a
 356 major gradient in soil fertility and light. DCA2 sam-
 357 ple scores had the highest correlation with Ellenberg
 358 (*L*) indicating a light gradient but also high correlation

359 to soil fertility. DCA3 sample scores had a high cor-
 360 relation with variables related to soil moisture-related
 361 variables (Ellenberg *F* value, *C* total). There was a
 362 higher species turnover among the abundant species
 363 on conventional farms than on organic farms (Fig. 2),
 364 and organic plot were positioned toward the higher
 365 fertility end of the gradient.

366 Total inertia was 8.28 and the variance explained
 367 by all explanatory variables was 2.03 (=25%). Fertili-
 368 ty, location, farm type, four soil variables and bank

Table 5

Correlation coefficients between explanatory variables and sample ordination score for DCA1–DCA3 and species richness

Explanatory variables	DCA1	DCA2	DCA3	Species richness
DCA1	1.00	0.05	0.11	0.003
DCA2	0.05	1.00	-0.07	0.12
DCA3	0.11	-0.07	1.00	0.03
pH	-0.13	-0.13	0.15	0.10
Conductivity	0.02	-0.09	-0.19	0.04
Water content	0.11	0.20	-0.38	0.17
Light	0.38	-0.18	0.27	0.12
Bank	0.01	0.29	0.09	-0.12
Exposition	0.23	0.13	-0.11	0.04
Total P	-0.32	-0.36	-0.07	0.17
Total N	0.29	0.19	-0.30	0.12
Loss on ignition	0.16	0.20	-0.33	0.09
Coarse sand	0.42	0.22	0.29	-0.34
Coarse silt	-0.15	-0.02	-0.26	0.12
Clay	-0.27	0.03	-0.27	0.19
Silt	-0.28	-0.27	-0.04	0.42
Total C	0.21	0.20	-0.36	0.10
Ellenberg (<i>L</i>)	-0.09	-0.53	0.38	0.09
Ellenberg (<i>T</i>)	0.24	-0.31	0.35	0.15
Ellenberg (<i>F</i>)	-0.14	0.15	-0.50	0.20
Ellenberg (<i>R</i>)	-0.35	-0.29	-0.38	0.34
Ellenberg (<i>N</i>)	-0.57	-0.39	-0.12	0.22

Values above 0.3 are represented in bold.

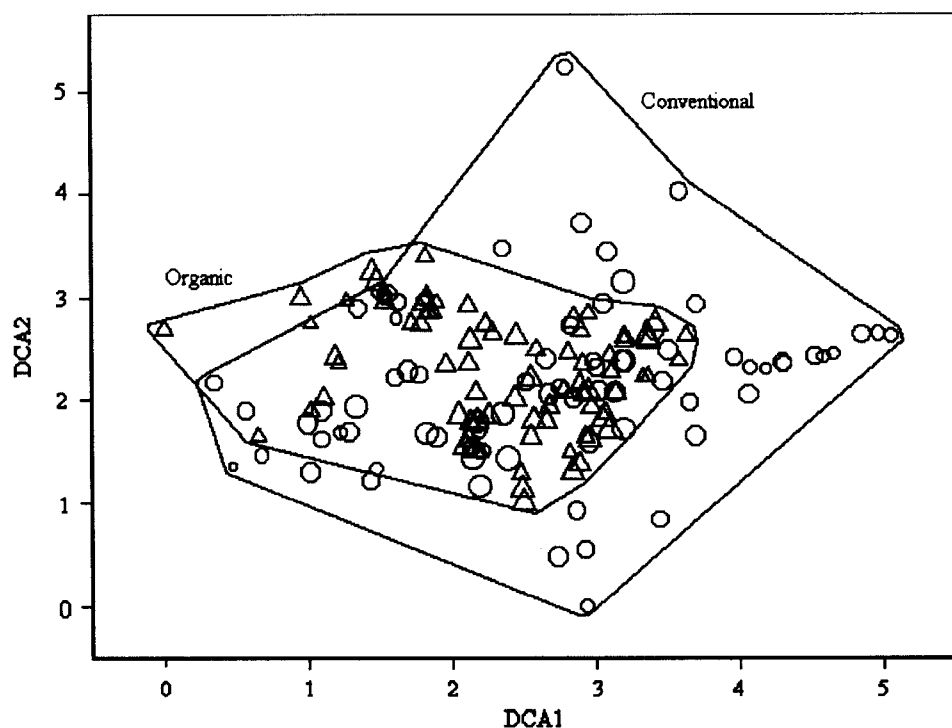


Fig. 2. Ordination diagram with 156 sampling plots along the two most important gradients, DCA1 and DCA2. ○: conventional hedgerow; △: organic hedgerow; polygons: surface covered by the each management types. Sizes of circles and triangles reflecting number of species in the sample plot.

369 height explained a significant fraction of the variation
 370 in species composition (Table 1). Soil phosphorous
 371 content was the variable that explained most of the
 372 variation (12%). Location of the sampling plots (in-
 373 side the hedgerow or outside in the open vegetation)
 374 was second (10.4%), farming type being only the fifth
 375 most important variable explaining 9.4% of the varia-
 376 tion (Table 1).

377 4. Discussion

378 Both null hypotheses have to be rejected because
 379 farming type significantly affected alpha, beta and
 380 gamma plant species diversity in the hedge bottom
 381 vegetation. Farming type explained a significant part
 382 of the variation in the species composition.

383 Farming type had a highly significant influence on
 384 the hedge bottom vegetation. A significantly higher
 385 species diversity was found in hedgerows without pes-
 386 ticide impact and adjacent to organic fields compared

387 to hedgerows on conventional farms. This difference in
 388 species diversity was significant in alpha and gamma
 389 diversity. The difference was even more obvious when
 390 only samples from hedgerow edges were compared.
 391 Other studies also found significantly higher species
 392 diversity in hedge bottom vegetation associated with
 393 more extensive farming systems (Hegarty et al., 1994;
 394 Boutin and Jobin, 1998; French and Cummins, 2001).
 395 However, the highly significant difference in hedge
 396 bottom vegetation between conventional and organic
 397 farming in this study had not been observed before. In
 398 many cases the effect of management can be difficult
 399 to interpret because hedges often have different land
 400 use on either side (Mercer et al., 1999) but that was
 401 not the case in this study.

402 Bank height and orientation differed significantly
 403 between farming types. Species diversity also corre-
 404 lated positively to bank height in Mercer et al. (1999)
 405 study. In our study there was only 0.06 m difference
 406 in bank heights between farming types, no difference
 407 in median values, there was a small negative cor-

408 relation between species richness and bank height,
409 and the highest mean value of banks was found in
410 conventional hedgerows. The significant difference
411 in orientation was unlikely to explain differences in
412 species richness as already recorded by other Euro-
413 pean workers (Mercer et al., 1999; Marshall et al.,
414 2002).

415 All hedges were located in the same Main Ter-
416 minal Moraine, on the same soil (Madsen and Jensen,
417 1992) and with only minor differences in climatic con-
418 ditions (Lysgaard, 1979). Hedgerows had the same
419 width, age and measured soil variables did not dif-
420 fer between farm managements. The potential local
421 and regional plant species pool (Zobel, 1997; Zobel
422 et al., 1998) was assumed to be the same and dif-
423 ferences in species number must be explained by ei-
424 ther extinction or immigration or both (MacArthur and
425 Wilson, 1967; Hanski and Gilpin, 1997). Higher ex-
426 tinction on conventional farms could be due to pesti-
427 cide drift but also from the phase of hedgerow estab-
428 lishment. Herbicides were applied to the conventional
429 hedgerows 2 or 3 years after establishment, whereas
430 organic farmers only used non-chemical weed con-
431 trol, the two weed control strategies favouring differ-
432 ent plant species (Kleijn and Snoeiijing, 1997). It is
433 impossible to tell whether it is difference in the es-
434 tablishment of organic and conventional hedgerows
435 or the following 10–14 years differences in farming
436 type that is most important. However, the use of her-
437 bicides was the main difference during both establish-
438 ment and adolescence. Field margins and non-target
439 plants could receive up to 20–25% of the applied pes-
440 ticide field dosage (De Snoo and de Wit, 1998; De
441 Snoo, 1999; Boutin et al., 2001), which may influ-
442 ence diversity and cover of non-target plant of field
443 margins (Marrs et al., 1989, 1991; Marrs and Frost,
444 1997; Hald et al., 1994; Jobin et al., 1997; Kleijn and
445 Snoeiijing, 1997; De Snoo and van der Poll, 1999).
446 Spray drift also influences the performance, survival
447 and reproductive potential of some non-crop species
448 (Fletcher et al., 1996). On the other hand, some studies
449 found that reduced pesticide drift were of minor im-
450 portance for the boundary flora (Hald, 1988; Marshall,
451 1992) which can be explained by the inertia of estab-
452 lished plant communities (Milchunas and Lauenroth,
453 1995).

454 Organic fields often have higher weed species
455 density and diversity (Hald, 1999; Rydberg and

Milberg, 2000) and the potential immigration rate 456
of weeds from the fields is likely to be higher com- 457
pared to conventional hedgerows. This is consistent 458
with the higher richness of weedy species recorded 459
in the hedge bottom of the organic farms. Despite 460
the barriers of movement encountered by many plant 461
species in modern fragmented landscapes (Primack 462
and Miao, 1992; Wiens, 2001) there is a high prob- 463
ability that non-weedy species have dispersed from 464
nearby semi-natural habitats. 465

466 There were few forest species and no difference
467 between farming type could be detected. Fertilising
468 can affect the plant species diversity (Marshall et al.,
469 2002), but is unlikely to explain the differences be-
470 tween farming types. There was no significant differ-
471 ence between farming types in either measured soil
472 fertility or calibrated Ellenberg values of fertility. A
473 difference in available *P* and *N* fractions between con-
474 ventional and organic farms cannot be excluded to-
475 tally, but Mäder et al. (2002) suggested that with equal
476 total phosphorous contents a higher available *P* is ex-
477 pected on organic farms due to higher microbial ac-
478 tivity.

479 Some studies indicate relations between landscape
480 ecological features and species diversity (Dzwonko
481 and Loster, 1988; Skånes, 1990) and composition
482 (Dzwonko, 1993; Dzwonko and Loster, 1997; Graae,
483 2000; Le Coeur et al., 1997, 2002). In this study,
484 hedgerows were positioned close to each other in the
485 same type of landscape and differences in species di-
486 versity can hardly be explained by differences in the
487 landscape context.

488 Disturbance can also be important in terms of
489 species diversity (Grime, 1979; Eriksson, 1997), be-
490 cause a closed sward may prevent recruitment of
491 species. Hedgerows near cereal fields could contain
492 more ruderal species than hedgerows close to grass
493 ley. In this study, more conventional hedgerows were
494 close to cereal fields than organic hedgerows (eight
495 versus six). Therefore, differences in disturbance
496 regime are not likely to explain the differences in
497 species diversity.

498 This study included more conventional than or-
499 ganic farms (eight versus five) which may result in
500 a higher probability of finding different local species
501 pools. Conventional farming may include a more di-
502 verse group of farming practices than organic farms
503 but it is more likely that the 25% lower alpha diver-

504 sity in conventional hedgerows resulted in lower beta
505 diversity.

506 The higher difference in species richness between
507 margin and centre plots in conventional than organic
508 hedgerows indicated a stronger agricultural impact
509 from the field to the centre of the hedgerow.

510 The higher frequency of species occurring in semi-
511 natural habitats and the hosting of a protected species
512 (*E. helleborine*) imply that a slightly higher natural-
513 ness (*sensu*, Angermeier and Karr, 1994; Anderson,
514 1991) occurred in hedge bottoms associated with or-
515 ganic farming systems. A higher number of species
516 could also act as a potential buffer against environmen-
517 tal fluctuations as suggested by Loreau et al. (2001)
518 and changing landscapes. On this background it can-
519 not be excluded that organic farming contribute to a
520 higher biological integrity and a more sustainable de-
521 velopment of the landscape.

522 Acknowledgements

523 We thank the farmers for letting us conduct the study
524 on their farms. Many thanks to Simon Smart, Rasmus
525 Ejrnæs and two anonymous referees for criticising the
526 manuscript. The Danish EPA (Miljøstyrelsen) partly
527 financed this study and contributed with valuable sta-
528 tistical discussions. Thanks to Anfred Pedersen for
529 identifying subspecies of *Rubus fruticosus*.

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