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To cite this article: Anna Kövendi-Jakó, Sára Márialigeti, András Bidló & Péter Ódor (2016) Environmental drivers of the bryophyte propagule bank and its comparison with forest-floor assemblage in Central European temperate mixed forests, *Journal of Bryology*, 38:2, 118-126, DOI: [10.1080/03736687.2015.1115804](https://doi.org/10.1080/03736687.2015.1115804)

To link to this article: <http://dx.doi.org/10.1080/03736687.2015.1115804>



Published online: 10 Feb 2016.



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Environmental drivers of the bryophyte propagule bank and its comparison with forest-floor assemblage in Central European temperate mixed forests

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Species richness, composition and abundance of the bryophyte diaspore bank of Central European temperate mixed forests were compared with the forest-floor bryophyte assemblage. The impact of environmental variables and anthropogenic disturbances, including tree species composition, stand structure, microclimate, light conditions, soil and litter properties, management history, and landscape properties, potentially influencing bryophyte diaspore bank assemblages were explored. Thirty-four, 70–100 years old mixed stands with differing tree species composition were examined in the Őrség National Park, Western Hungary. The diaspore bank was studied by soil collection and cultivation, and data were analysed by multivariate methods. Contrary to the forest-floor bryophyte assemblage, where substrate availability, tree species composition and stand structure were the most influential environmental variables, the composition and abundance of the diaspore bank was mainly affected by site conditions (microclimate, litter and soil properties). Species richness of the bryophyte diaspore bank was lower than that of the forest-floor bryophyte assemblage. Short-lived mosses (colonists, short-lived shuttles) were dominant in the diaspore bank, as opposed to the forest-floor bryophyte community, where perennial mosses dominated. In the studied forests, the importance of the bryophyte diaspore bank was relatively low in the regeneration and maintenance of the forest-floor bryophyte vegetation.

Keywords: Diaspore bank, Ordination, Species composition, Species richness

Introduction

Reservoirs of dormant reproductive propagules are known from several groups, such as phanerogams (seed bank), ferns, bryophytes (diaspore bank), dinoflagellates and copepods. Their role is increasingly appreciated in nature conservation, restoration, life history analysis and population genetics. The diaspore bank of bryophytes is less well known than the soil seed bank of phanerogams (During, 1997; Csontos, 2001). The soil contains a “bank” of bryophyte diaspores, including “a number of dormant, ungerminated but viable spores and/or asexual propagules: rhizoidal or protonemal gemmae, tubers, and resting buds on stem fragments” (During, 2001). The different types of propagules have been generally thought to have different functions: the spores are mainly important for long-distance dispersal while asexual reproduction maintains local populations.

The lifetime of spores is longer, their dispersal ability is better and their germination capability is worse than that of asexual diaspores (During, 1979). The production of spores is usually strongly seasonally determined (Hock *et al.*, 2004), while the production of asexual diaspores is usually much less seasonal (During, 1997, 2001). In contrast to the seed bank of phanerogams, the bryophyte diaspore bank is dominated by asexual propagules (During, 1997, 2001; Longton, 1997; Söderström & Herben, 1997; Laaka-Lindberg, 1999; Rydin, 2009). Bryophytes are classified into life strategy types, based on the life span, reproductive effort of the gametophyte, and size and abundance of spores (During, 1979, 1992). Usually bryophyte species are not all represented equally well in the diaspore bank. Pleurocarpous, perennial bryophytes that dominate the above-ground vegetation tend to be underrepresented in the diaspore bank, because they have low reproductive effort and long life span in the forest-floor assemblages. By contrast,

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acrocarpous colonists, and annual and short-lived shuttle species are dominant in the soil diaspore bank even in the vegetation types where the above-ground vegetation is dominated by perennial pleurocarpous bryophytes. These bryophyte species have persistent spores and high sexual and asexual reproductive effort, short life span in the forest-floor vegetation and they have an important role in the regeneration of forest-floor bryophyte vegetation after disturbance (During, 1979, 1992, 1997, 2001; Vanderpooren & Goffinet, 2009). Many species of the propagule bank are not well represented in the vegetation; even scientifically new taxa have been described from diaspore bank, such as the species *Neophoenix matoposensis* and *Ephemerum* spp. (Zander & During, 1999).

Most studies have demonstrated that the diaspore bank has similar or greater diversity than the above-ground vegetation (Jonsson, 1993; Rydgren & Hestmark, 1997; Hock *et al.*, 2008). In addition to the aforementioned importance in bryophyte diversity, the role of the bryophyte propagule bank is diverse: ensuring the survival of bryophytes under unfavourable conditions and after disturbance, facilitating regeneration and colonization, and regulating population size (Van Tooren *et al.*, 1990; Jonsson, 1993; Lloret, 1994; During, 1995, 1997; Hock *et al.*, 2008; Caners *et al.*, 2009; Vanderpoorten & Goffinet, 2009). While for phanerogams many types of dormancy and dormancy-breaking mechanisms have been described, in the case of bryophyte propagules, knowledge is sparse (Sundberg & Rydin, 2000; During, 2001; Hock *et al.*, 2004).

Most important drivers of species composition, structure and spatial pattern of bryophyte assemblages are different for the forest-floor vegetation and propagule bank, but forest-floor vegetation has been more intensively studied. In temperate and boreal forests, the most important drivers for forest-floor bryophytes are mainly microclimate (Mills & Macdonald, 2005; Hylander & Dynesius, 2006), soil conditions (Hylander & Dynesius, 2006) and availability of appropriate substrates (Schofield, 1985; Ódor & Standovár, 2002; Ódor & van Hees, 2004; Wiklund & Rydin, 2004). Márialigeti *et al.* (2009) demonstrated that the litter of deciduous trees has a negative effect on the occurrence of forest floor bryophytes, which are restricted to open mineral soil and logs. Stand structure heterogeneity, tree species richness and the presence of a regeneration (shrub) layer considerably increases bryophyte cover and diversity (Aude & Lawesson, 1998; Márialigeti *et al.*, 2009).

In contrast to forest-floor bryophytes, the relationship between forest site conditions and the bryophyte diaspore bank has hardly been studied (Caners *et al.*, 2009). Comparative studies between forest-floor

bryophytes and diaspore banks in forests are also sparse (During *et al.*, 1987; Jonsson, 1993; Rydgren & Hestmark, 1997; Hebrard, 2001; Rydgren *et al.*, 2004; Szűcs, 2013; Iglesias *et al.*, 2015). Our investigation aimed to explore the environmental drivers of the diversity and composition of the bryophyte diaspore bank of temperate mixed forests, and to compare it with the forest-floor bryophyte assemblage. The study covered a wide range of potentially influential environmental variables (including tree species composition, stand structure, microclimate, light conditions, soil and litter properties, management history and landscape properties) in order to select the most important ones for diaspore bank composition and richness. In addition, the species richness and species composition of the diaspore bank and the forest-floor bryophyte vegetation were compared.

Materials and Methods

Study area

The study was located in the Órség National Park, Western Hungary (46°51–55'N and 16°07–23'W). The landscape consists of hills and wide valleys, 250–350 m above sea level. The mean annual precipitation is 700–800 mm and mean yearly temperature is 9.0–9.5°C (Gyöngyössi, 2008). The superficial deposits are composed of Pleistocene sediments including loess and alluvial gravel. The soil is acidic (pH 4.5–4.7) and nutrient poor; the most common soil type on hills is pseudogleyic and lessivaged brown forest soil (Luvisols), while in the valleys mire and meadow soils (Gleysols) can be found (Szodfridt, 1969).

Órség is a unique part of Hungary because different combinations of tree species appear under similar site conditions. Many tree species are found with relatively high abundance and various proportions in the stands, the dominant ones being beech (*Fagus sylvatica*), sessile and pedunculate oak (*Quercus petraea* and *Quercus robur*), hornbeam (*Carpinus betulus*), Scotch pine (*Pinus sylvestris*) and Norway spruce (*Picea abies*). The proportion of different associated species (such as *Betula pendula*, *Populus tremula*, *Castanea sativa*, *Prunus avium*) is also considerable. Most of the forest stands are mixed, with relatively high tree species diversity. Differences in species composition are the result of climate, and historical land use patterns.

This region of Hungary has been influenced by agriculture for centuries. Since the twelfth century extensive farming has decreased forest cover and resulted in a shifting mosaic of land use types. Litter collection in forests and a special form of tillage, ridging, changed the vegetation, and also resulted in the acidification of the soil and increased soil erosion. These processes are advantageous to Scotch pine

regeneration, the establishment of a high terricolous bryophyte cover (*Pleurozium schreberi*, *Scleropodium purum*, *Dicranum polysetum*, *Polytrichum formosum*), and acidophytic shrub and herbaceous species (Ódor *et al.*, 2002; Tímár *et al.*, 2002). From the nineteenth century extensive farming decreased; the former arable lands reverted to forests, dominated by Scotch pine and other pioneer tree species. Two types of forest management are used nowadays: in private forests, a stem selection system is followed resulting in uneven aged stands; in state forests, a shelterwood management system is used, creating even aged stands with a rotation period of 70–100 years. After the stabilization of land use types, the species composition of the managed forests is changing, the dominance of deciduous tree species (beech, oaks, hornbeam) is increasing, while that of Scots pine and pioneer trees is decreasing (Tímár *et al.*, 2002).

Data collection

Thirty-five forest stands (2–15 ha each) were selected for this study by stratified random sampling from the stand structural database of the Hungarian National Forest Service. Criteria of site selection were as follows: all stands were 70–100 years old, located on level areas (slopes were less than 10°), and not directly influenced by water. The 293 stands meeting these criteria were grouped according to tree species composition representing different combinations of the main tree species (beech, sessile and pedunculate oak, Scots pine). The main groups were monodominant stands of one species (cover higher than 80%) and mixed forests dominated by two or three of the aforementioned species. The 35 stands were randomly selected from these groups. The selected stands represent old growth forests, covering the different common tree species combinations in the area. The abundance of forest-floor bryophyte species (including species occurring on lying dead wood) was expressed in cover, which was estimated in dm² within each 30 m × 30 m plot. The total abundance of the terrestrial bryophyte layer per plot was expressed as the sum of the cover of the individual species.

Three bryophyte diaspore bank samples were collected in each 30 m × 30 m plot in November 2009. Soil cubes with 7 cm side length were taken out after the litter and other organic materials were removed from the surface. The soil cubes were put into paper bags and transported to the laboratory for cultivation. Altogether 105 sampling units were collected (35 plots, three subunits per plot).

During cultivation, a thin layer of the soil sampled (0.5–1.5 cm deep) was spread out in closed, transparent boxes (with 10 cm radius) on a layer of *ca* 2–4 cm of autoclaved perlite, moistened with deionized water. Three control boxes containing autoclaved perlite

layer with a thin autoclaved soil layer were also included for detecting air-borne propagule contamination. The samples were cultivated in greenhouse for 3 months as in Hock *et al.* (2004), finishing at the end of January 2010. Cultivation was at room temperature (18–21°C), with natural (from 8 to 16 hours) and artificial (from 16 to 18 hours) light conditions (1200–2200 lx).

The abundance of the diaspore bank species within the sampling units was expressed by local frequency values. A frame 7 cm × 7 cm divided into 1 cm² sized squares by a grid was put into the cultivation box. Presence/absence of species was recorded within the squares. The maximum local frequency value of a species was 147 (3 × 49) within a plot. Abundance of the whole diaspore bank was expressed as the sum of the local frequencies of the individual species. Voucher specimens are deposited in the private herbarium of the last author. The identification of the young, atypical specimens was sometimes complicated (During, 1997). *Sphagnum* spp. and *Ditrichum* spp. (mainly *D. pusillum*, *D. cylindricum* and *D. pallidum*) were identified only at genus level. Nomenclature follows Erzberger and Papp (2004) for bryophytes and Tutin *et al.* (1964–1993) for vascular plants.

Fifty-two environmental variables were recorded for analysis (Table 1). Variables related to tree species composition and tree size (species identity, diameter, height) were based on measurements of tree individuals larger than 5 cm diameter at breast height (DBH) within the plots. Number of saplings (including trees and shrubs) thinner than 5 cm DBH but taller than 0.5 m were counted and their density (unit per hectare) was calculated. For dead wood volume calculation, only individuals thicker than 5 cm were included (snags and logs separately). Cover of understory vegetation (including herbs and seedlings shorter than 0.5 m), bryophytes and different substrate types (mineral soil, deadwood and litter) were estimated (on plots of 30 m × 30 m). Canopy openness was measured with a spherical densitometer, and relative diffuse light with LAI-2000 Plant Canopy Analyser instruments; in both cases, the measurements were carried out at 36 points within the plots at 1.3 m height (Tinya *et al.*, 2009).

Microclimate (air humidity and temperature) was measured in the middle of the plots at 1.3 m height using Voltcraft DL-120 TH data loggers in 24-hour measurement durations, eight times during two years. The microclimate of the forest stands in general, rather than that of the bryophyte communities (*sensu* Király *et al.*, 2013) was recorded. Characteristics of litter and soil were surveyed in five sampling units per plot. Litter was collected from 30 cm × 30 cm areas, but only the upper 10 cm layer was studied. The litter properties analysed were pH, weight, composition of litter

Table 1 Potential explanatory variables used to characterize the ecological conditions of the analysed forest stands

<i>Tree species composition</i>	
Tree species richness	
Tree species Shannon diversity	
Relative volume of tree species (beech, hornbeam, oaks, pine, spruce, mixed tree species) (%)	
<i>Stand structure</i>	
Size distribution of trees (mean, coefficient of variation and diversity of DBH; cm)	
Density of large trees (DBH > 50 cm) (stems/ha)	
Sapling density (stems/ha)	
Understorey cover (m ² /ha)	
Timber volume (m ³ /ha)	
Dead wood volume (snag and log separately; m ³ /ha)	
<i>Substrate types</i>	
Cover of mineral soil, litter and dead wood (m ² /ha)	
<i>Characteristics of litter</i>	
Proportion of coniferous litter (%)	
Litter weight (on an area of 30 × 30 cm; g)	
<i>Light conditions</i>	
Relative diffuse light (mean, coefficient of variation, %)	
Canopy openness (mean, standard deviation; %)	
<i>Landscape variables</i>	
Proportion of landcover types in the surroundings of plots (radius = 300 m; old beech, oak, Scotch pine, Norway spruce and mixed forests, young forests, non-forested areas; %)	
<i>Management history</i>	
Proportion of cultivation types in the surroundings of plots (radius = 300 m) in 1853 (forest, meadow, arable land; %)	
Cultivation types of the plots in 1853 (forest, meadow, arable land; %)	
<i>Characteristics of forest site</i>	
Litter properties (pH, weight (g), composition, carbon and nitrogen content; %)	
Soil properties (pH, acidity, texture (%), carbon (%), nitrogen (%), phosphorous and potassium content; mg/100 g)	
<i>Microclimate</i>	
Humidity (mean and the coefficient of variation; %)	
Temperature (mean and coefficient of variation; °C)	

types (deciduous, coniferous, decayed), carbon and nitrogen content. We also considered the following mineral soil properties: pH, acidity, texture, carbon, nitrogen, phosphorous and potassium content (Juhász et al., 2011). The proportion of land-cover types were estimated within circles of 300 m radius around the plots by remote sensing based on aerial photos, maps and data from the Hungarian Natural Forest Service (Márialigeti et al., 2009). Historical variables were generated based on the map of the Second Military Survey of the Habsburg Empire from 1853 (Arcanum, 2006). The management type of the plots and the proportion of these within a circle of 300 m radius were calculated (Table 1; Király & Ódor, 2010).

Data analysis

The species richness, abundance and species composition of the diaspore bank and the forest-floor bryophyte assemblages were compared both at plot level (alpha diversity) and whole sample species richness (gamma diversity). Because of non-normal distribution and non-equal variances, plot level species richness was compared by Mood median test (Zar, 1999). Correlation of plot level species richness and plot level abundance was calculated between the two assemblages. The species composition was compared based

on frequency data (based on occurrences in 35 plots) and the abundance data of the species. The relationship between the composition of the two assemblages was expressed as the matrix correlation of their distance matrices (Mantel-test, Legendre & Legendre, 1998). Euclidean distance matrices between the plots were calculated for the two assemblages, based on the abundance values of the species (for diaspore bank: local frequency data; for forest-floor vegetation cover data were used). The significance of the matrix correlation coefficient was tested by Monte-Carlo simulation, and the number of permutations was 1000. The distribution of life strategy types (During, 1992) of the two assemblages was compared by chi-square analysis of contingency tables (Zar, 1999).

Relationships between the bryophyte species composition of the diaspore bank and potential explanatory variables were explored by indirect (principal component analysis, PCA and detrended correspondence analysis, DCA) and direct (redundancy analysis, RDA) ordination (Legendre & Legendre, 1998; Podani, 2000). Species data were centred and logarithmically transformed. Bryophyte species with lower frequency than four (occurred in less than five plots out of 35) were omitted from the analysis. As the gradient length of the DCA axes was lower than 1.5

standard deviation unit, RDA was used for direct ordination. This method supposes an approximately linear relationship between the species abundance and the explanatory variables. The environmental factors of the redundancy analysis were selected by manual forward selection. The effect of factors was tested by F-statistics via Monte-Carlo simulation, the number of permutations was 499, and the threshold significance level was 0.05 (Ter Braak & Smilauer, 2002; Lepš & Šmilauer, 2003). The significance of all canonical axes was investigated similarly to the explanatory variables. Because of its significant effect, the longitudinal geographical coordinate was used as a covariate in the final ordination.

The relationship between the abundance of diaspore bank (dependent variable) and the environmental (explanatory) variables was tested by linear modelling, using Gaussian error structure without transformation (Faraway, 2005). As the longitudinal geographical coordinate had a significant effect, it was used as a covariable within a partial multiple regression framework (Legendre & Legendre, 1998). Before modelling, preliminary selection and data exploration were performed. Only those explanatory variables were included in the linear model selection that (1) correlated significantly with the dependent variables; (2) had homogenous scatter plots with them; and (3) their inter-correlations with the other explanatory variables were low (the absolute values of the correlation coefficients were lower than 0.40). The minimal adequate model was built by backward elimination, using deviance analysis with F-test (ANOVA). The model selection was also checked by likelihood methods using the Akaike Information Criterion (Faraway, 2005). The normality and variance homogeneity of the residuals of the final model were checked. Multivariate analyses were carried out with Canoco for Windows 4.5 (Ter Braak & Smilauer, 2002), whereas the rest of the analyses were carried out in the R 2.14.0 statistical environment (The R Development Core Team, 2011), using the vegan package for the Mantel-test (Oksanen et al., 2011).

Results

Comparison of diaspore bank and forest-floor assemblages

The diversity of the diaspore bank was lower than that of the forest-floor bryophyte assemblage. Only 13 bryophyte species occurred in the diaspore bank, while the whole sample of the forest-floor assemblage contained 70 species (Supplementary Material 1, Table S1). The mean species richness of plots was 5.80 (median 6) for the diaspore bank and 19.20 (median 18) for the forest-floor assemblage; the median of the two variables significantly differed

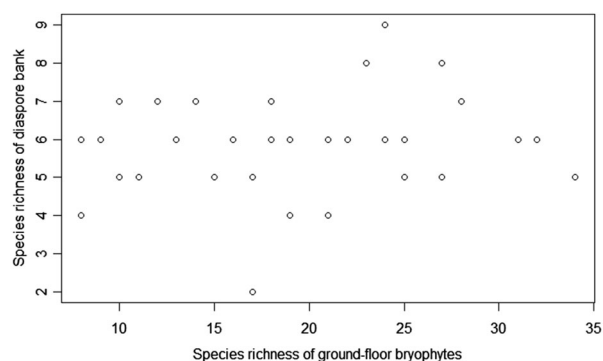


Figure 1 Relationships between forest-floor and diaspore bank assemblages according to species richness (species number per plot).

(Mood median test, $P < 0.001$). Neither the species richness (Figure 1, $r = 0.17$, $P = 0.17$) nor the abundance ($r = 0.07$, $P = 0.67$) of forest-floor and diaspore bank assemblages correlated. The composition of the life strategy types based on species abundance was different between the two assemblages (Figure 2, test of contingency tables, $\text{Chi}^2 = 77375$, $\text{df} = 3$, $P < 0.001$). The diaspore bank was dominated by colonists, the forest-floor assemblage by perennials. The only exception was the perennial *Atrichum undulatum*, which was found frequently in both assemblages.

The species composition of the assemblages was significantly different, and the distance matrices of the plots derived from the two assemblages were significantly not correlated (Mantel-test, $r = 0.151$; $P = 0.095$). Many species occurred in the forest-floor vegetation but were absent from the diaspore bank, resulting in the lower species richness of the second assemblage (Supplementary Material 1, Table S1). The overlap in the species composition was low. Only *Atrichum undulatum*, *Dicranella heteromalla* and *Pohlia nutans* were abundant in both assemblages. However, many species (*Dicranella rufescens*, *Philonotis arnellii*, *Physcomitrium pyriforme*, *Sphagnum* spp. and *Trichostomum*

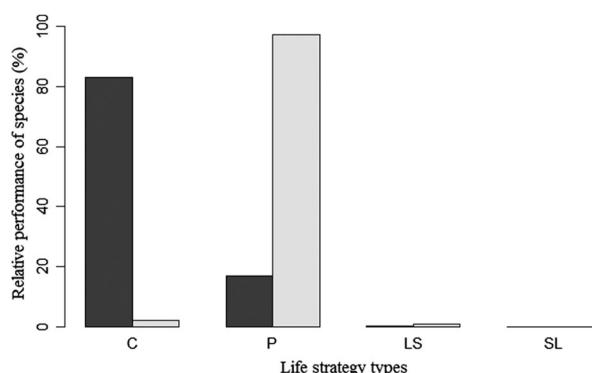


Figure 2 Distribution of the life strategy types in diaspore bank (black) and forest-floor (gray) assemblages based on the relative abundance of species. Strategy types: C – colonist; P – perennial; LS – long lived shuttle; SL – short lived shuttle.

brachydontium) occurred only in the diaspore bank, while *Bryum rubens* and *Ditrichum* spp. were dominant in the diaspore bank but rare in the forest-floor assemblage. Many abundant terricolous-opportunistic species (e.g. *Brachythecium rutabulum*, *Dicranum scoparium*, *Hypnum cupressiforme*, *Polytrichum formosum*) and epiphytic-epixylic (*Dicranum montanum*, *Frullania dilatata*, *Lophocolea heterophylla*, *Platygyrium repens*, *Plagiothecium* spp., *Radula complanata*) species of the forest-floor vegetation were absent from the diaspore bank. Some abundant species of the forest-floor assemblage occurred with very low local frequency in the diaspore bank (e.g. *Brachythecium velutinum*, *Herzogiella seligeri* and *Rhizomnium punctatum*). There was no germination in the three autoclaved control boxes included for detecting air-borne propagule contamination.

Multivariate analysis of the diaspore bank

The first and second PCA axes explained 43% and 22% of species variance, respectively. In contrast to species, the appearance of the plots along the PCA axes was not grouped (Figure 3). *Atrichum undulatum*, *Dicranella heteromalla* and *Pohlia nutans* had negative values along the first PCA axis, while *Bryum rubens*, *Dicranum rufescens* and *Ditrichum* spp. were separated based on the second PCA axis (they had negative values on it).

The canonical axes of RDA explained 45.8% of total variance (first axis 23.4%, second 14.6%, Figure 4). The effects of the first axis ($F = 8.27$, $P = 0.002$) and all canonical axes ($F = 3.81$, $P = 0.002$)

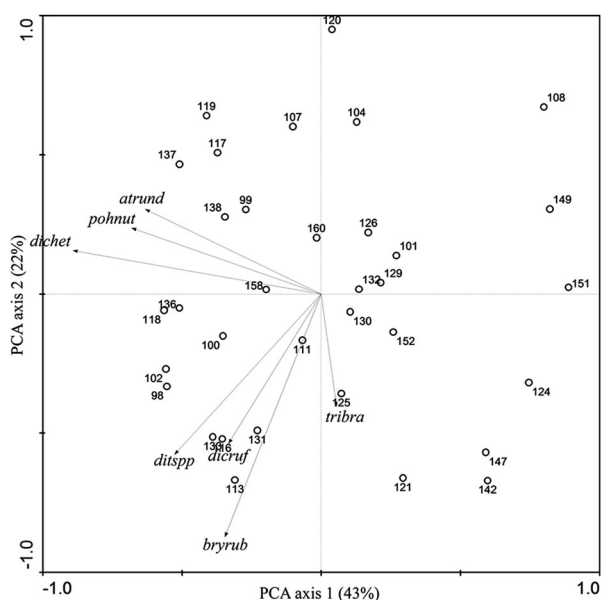


Figure 3 Biplot of the principal component analysis (PCA) of the diaspore bank. The first axis explained 43%, the second 22% of the species variance. Species are marked by arrows and abbreviations, and the plots by open circles and numbers. Species abbreviation consists of the first three letters of the genus and species names (Supplementary Material 1, Table S1).

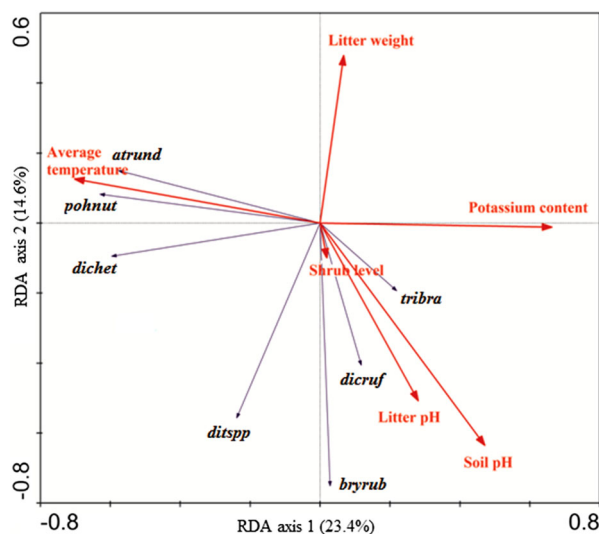


Figure 4 Species and environmental variables biplot of the redundancy analysis (RDA). The first axis explained 23%, the second 15% of the species variance. Species abbreviation consists of the first three letters of the genus and species names (Supplementary Material 1, Table S1). Explanatory variables are represented with red arrows and letters, for description see Table 1.

Table 2 Names (variables), explained variance (variance%), F values (F) and significance (P) of the selected explanatory variables of the redundancy analysis

Variables	Variance (%)	F	P
Average temperature	13	4.68	0.004
Soil pH	9	3.57	0.016
Soil potassium content	7	2.96	0.032
Litter weight	6	2.70	0.026
Litter pH	6	2.87	0.024
Shrub density	5	2.46	0.038

were significant. Average temperature, soil pH, soil potassium content, litter weight, litter pH and shrub density were significant explanatory variables in the final RDA model (Table 2). The first axis correlated positively with soil potassium content and negatively with average temperature. The second axis had positive correlation with litter weight and negative with shrub density, litter pH and soil pH. Three dominant acrocarpous species, *Atrichum undulatum*, *Dicranella heteromalla* and *Pohlia nutans* had positive correlation with average temperature and negative correlation with soil potassium content. Presence of *Bryum rubens*, *Dicranum rufescens* and *Ditrichum* spp. were affected mainly positively by shrub density, litter pH and soil pH, and they were negatively related to litter weight.

Based on the multiple regression model ($R^2 = 0.23$, $F_{3,31} = 4.35$, $P = 0.011$), the abundance of the diaspore bank was positively related to the cover of the understory and air temperature, while it was negatively related to soil acidity (Table 3).

Table 3 Importance of the explanatory variables in the regression model of the diaspore bank abundance

Variables	Estimate	Variance (%)	F	P
Understory cover	46.78	11.61	5.11	0.031
Acidity	-40.25	8.29	3.65	0.065
Temperature	42.05	9.73	4.29	0.047

Discussion

Comparison of diaspore bank and forest-assemblages

In this study, the direct comparison of forest-floor and diaspore bank species richness is difficult, because of the different sampling methodology: ground floor species richness was surveyed in the whole 900 m² area, while the diaspore bank was analysed from 3 ca 50 cm³ soil cubes. The sampling intensity for diaspore bank was too low to estimate species richness for the whole (900 m²) area. It is also possible that not all potentially viable propagules germinated during our experiment. However, the compositional differences between diaspore bank subunits (expressed as Euclidean distance) were significantly lower for within plot than for between plot subunits (Supplementary Material 2). We interpret that the beta diversity of subunits within a plot is relatively low. Although the species richness of the diaspore bank was probably underestimated because of methodological limitations, the species richness difference between the diaspore bank and forest-floor vegetation was so high (13 versus 70 for the whole sample, 5.8 versus 19.2 for plot level) that it suggests important differences in total species richness between the diaspore bank and the forest-floor vegetation.

Contrary to our results, it has been shown that the diaspore bank can have similar or higher diversity than the actual forest-floor, both in temperate forests and grasslands (Jonsson, 1993; Hock, 2007; Hock et al., 2008; Vanderpoorten & Goffinet, 2009). In tropical rain forests, a similarly species poor diaspore bank was observed, because of the inhibitory effect of the litter layer (Maciel-Silva et al., 2012). The low diversity of the diaspore bank could be explained by the regular, long-term disturbance of the forest soil. Many stands were ploughed in the past, and the soil surface of the regenerated forests has been regularly disturbed by forestry activities. Chronic disturbance could deplete the diaspore bank (Jonsson, 1993; During, 1997), especially if its species are very rare in the forest-floor vegetation.

In our case, the low diversity of diaspore bank can partly be explained by the large compositional difference between the diaspore bank and the forest-floor assemblages. These two assemblages were different; only two species (*Atrichum undulatum* and *Dicranella heteromalla*) had considerable abundance in both

groups. Liverworts did not establish a diaspore bank in the studied temperate forests, which is confirmed by earlier results from boreal (Jonsson, 1993; Ross-Davis & Frego, 2004) and tropical regions (Maciel-Silva et al., 2012; but see Bisang et al., 2003; Caners et al., 2009). Some dominant species of the forest-floor vegetation that produce huge numbers of spores (e.g. *Hypnum cupressiforme* and *Polytrichum formosum*) were not present in the diaspore bank. The absence of *Polytrichum formosum* is a surprising result, because it was fairly abundantly present in several previous studies (During et al., 1987; Jonsson, 1993; Iglesias et al., 2015).

Short-lived colonist mosses were dominant in the diaspore bank, as opposed to the forest floor bryophyte community, where perennial mosses dominated. These results are confirmed by many previous studies (e.g. During, 1979, 1997; During & ter Horst, 1983; During & van Tooren, 1987). Szűcs (2013) noticed similar species in the diaspore bank of Hungarian beech forests, namely, *Pohlia nutans* was the most abundant species, and *Atrichum undulatum*, *Bryum* spp. and *Dicranella heteromalla* were also frequent.

Multivariate analysis of the diaspore bank

Atrichum undulatum, *Dicranella heteromalla* and *Pohlia nutans* preferred relatively warm, more open, nutrient poor stands, while *Bryum rubens*, *Dicranum rufescens* and *Ditrichum* spp. were associated with stands with relatively less acidic litter and soil conditions. Other studies focusing on the propagule bank also confirm that *Pohlia nutans* prefers relatively acidic soil conditions both in boreal (Rydgren et al., 2004; Caners et al., 2009) and mixed forests (Ross-Davis & Frego, 2004).

Litter accumulation could prevent the establishment of bryophytes in the study area, both in the diaspore bank and on forest floor (Márialigeti et al., 2009). A similar phenomenon was observed in other regions, both in wetlands (Peintinger & Bergamini, 2006) and forests (Dzwonko & Gawronski, 2002; Légraré et al., 2005; Startsev et al., 2008). Based on our results, both species composition and the abundance of the diaspore bank were positively influenced by higher air temperature. Relatively higher temperatures could accelerate the germination of the propagule bank after disturbance (During, 1979, 1997; Schofield, 1985; Risse, 1987; Wiklund & Rydin, 2004; Caners et al., 2009). Soil temperature and humidity can have a more direct effect on the propagule bank than the measured air temperature and humidity, but these variables are strongly related. Our findings that the potassium content had a negative effect, while soil and litter pH had a positive influence on the germination of the diaspore bank propagules have been supported by a number of studies (During, 1979, 1997;

Schofield, 1985; Risse, 1987; Wiklund & Rydin, 2004; Caners et al., 2009; Rydin, 2009; Maciel-Silva et al., 2012).

Light is a very important environmental variable for the forest bryophyte vegetation (Wiklund & Rydin, 2004; Kimmerer, 2005; Caners et al., 2009), but our results show that it was not a significant environmental variable for the diaspore bank. Light influences spore germination and gametophyte growth that could eventually influence competitive relations (During, 1979; Schofield, 1985; Vanderpoorten & Goffinet, 2009). The understory layer and shrub density had a positive effect on the propagule bank, similarly to our results for forest-floor bryophytes (Márialigeti et al., 2009). These factors could produce a favourable microclimate not only for the terrestrial bryophytes, but also for the diaspore bank after disturbance. Since light is not an influential environmental variable in our study, this positive effect of the microclimate may overwhelm the negative shading effect of the understory and shrub layers. Contrary to our results, herbaceous and bryophyte cover negatively influenced the germination of the diaspore bank in many cases (Virtanen et al., 2000; Peintinger & Bergamini, 2006). Similarly to Caners et al. (2009), landscape variables and management history were not directly influential on the propagule bank. The frequent disturbances in the past could have had an important effect on the deterioration of the propagule bank, but the effect of land use history is not traceable in the recent assemblages. The longitudinal geographical extent of the study had significant effect, but it is very difficult to relate it to direct environmental variables. Because its direct effect was not a target of this study we used it as a covariable during the analyses.

Summarizing, as opposed to the forest-floor bryophyte assemblages (Márialigeti et al., 2009), where substrate availability, tree species composition and stand forest structure were the most influential environmental variables, the diaspore bank was affected mainly by forest site conditions (microclimate, litter and soil properties). It is in accordance with the study of Caners et al. (2009) from boreal forests, in that light conditions, pH and potassium content were the most important factors affecting the diaspore bank.

Conclusions

Large differences were observed between the propagule bank and the forest-floor assemblages in respect of species richness, species composition and limiting environmental factors. Bryophyte species can regenerate from diaspore bank, by propagule dispersal or clonal expansion (Jonsson, 1993; Rydgren & Hestmark, 1997). In this study, we conclude that the role of the bryophyte diaspore bank is relatively low in the regeneration

and maintenance of the forest-floor bryophyte vegetation in the mixed forests of Central Europe. In the establishment, maintenance and regeneration of the forest-floor bryophyte vegetation, the most important processes are clonal expansion (regeneration) and colonization by airborne propagules. The diaspore bank can be an essential part of the life history for some short-lived species, but it is not important to the establishment of the dominant forest-floor species because they have different life histories, without the production of long-lived soil dwelling diaspores (During, 1979, 1997).

Acknowledgements

The authors thank Zsófia Hock for methodological advice, participants of the Örs-Erdő project for environmental data collection, and the Department of Plant Taxonomy, Ecology and Theoretical Biology for providing the infrastructure. This study was supported by the Hungarian Research Fund (OTKA 79158), the Directory of Órség National Park and the TÁMOP-4.2.2.D-15/1/KONV-2015-0023 project of the European Union and European Social Found. Péter Ódor is a grantee of the János Bolyai Scholarship of the Hungarian Academy of Sciences.

Supplemental data

Supplemental data for this article can be accessed 10.1080/03736687.2015.1115804.

Taxonomic Additions and Changes: Nil.

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