

Vascular plant species richness at the landscape scale: Patterns and processes

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Abbreviations

ATKIS	Authorative Topographic-Cartographic Information System, <i>Amtliches Topographisch-Kartographisches Informationssystem</i>
Az.	Reference number, <i>Aktenzeichen</i>
BfN	Federal Agency for Nature Conservation, <i>Bundesamt für Naturschutz</i>
BMU	Federal Ministry for the Environment, Nature Conservation, Building and Nuclear Safety, <i>Bundesministeriums für Umwelt, Naturschutz, Bau und Reaktorsicherheit</i>
BNatSchG	Federal Nature Conservation Act, <i>Bundesnaturschutzgesetz</i>
BÜK 50	Soil map, scale 1:50,000, <i>Bodenübersichtskarte 1:50,000</i>
CaCO₃	Calcium carbonate
CBD	Convention on Biological Diversity
COP	Convention of the Parties
DBU	German Federal Foundation for the Environment, <i>Deutsche Bundesstiftung Umwelt</i>
DEM	Digital Elevation Model
DHDN	<i>Deutsches Hauptdreiecksnetz</i> (coordinate system)
DLM	Digital Landscape Model
DWD	German Weather Service, <i>Deutscher Wetterdienst</i>
EEA	European Environment Agency
e. g.	For example [<i>lat. exempli gratia</i>]
EIV	Ellenberg Indicator Values
EU	European Union
FFH	Fauna-Flora-Habitat (Habitats Directive)
FG	Fragmentation geometry
FGSV	Road Research Institute, <i>Forschungsgesellschaft für Straßen- und Verkehrswesen</i>

GBIF	Global Biodiversity Information Facility
GIS	Geographic Information System
GLM	Generalised Linear Model
GSPC	Global Strategy for Plant Conservation
i. e.	that means [<i>lat. id est</i>]
IPA	Important Plant Areas
LCT	Land cover type
LGN	Lower Saxon Authority for Land-surveying and Geoinformation, <i>Landesvermessung und Geobasisinformation Niedersachsen</i>
m_{eff}	Effective Mesh Size Index
n/a	not available
NLFB	Lower Saxon Regional Authority for Mining, Power and Geology, <i>Landesamt für Bergbau, Energie und Geologie</i>
NLÖ	Lower Saxon Regional Authority of Ecology, <i>Niedersächsisches Landesamt für Ökologie</i>
NLWKN	Lower Saxon Water Management, Coastal Defence and Nature Conservation Agency, <i>Niedersächsisches Landesamt für Wasserwirtschaft, Küsten und Naturschutz</i>
NMUK	Lower Saxon Ministry for the Environment and Climate Protection, <i>Niedersächsischen Ministerium für Umwelt, Energie und Klimaschutz</i>
NR	Geographical regions, <i>naturräumliche Regionen</i>
NSG	Nature protection area, nature reserve, <i>Naturschutzgebiet</i>
NW-FVA	Northwest German Forest Research Station, <i>Nordwestdeutsche Forstliche Versuchsanstalt</i>
RL	Red List
UG	Study area, <i>Untersuchungsgebiet</i>
UNECE	United Nations Economic Commission for Europe
VEAC	Victorian Environmental Assessment Council
z. B.	German for <i>zum Beispiel</i> (for example)

Summary

The heterogeneous geographic distribution of species richness has fascinated and challenged biologists over centuries. As a result, a large number of possible hypotheses that aim at explaining these patterns have been proposed. Currently, the consensus is that multiple processes simultaneously affect species richness patterns, with the influence of each process differing between taxa, landscapes and periods in time. Generally, environmental conditions are assumed to be the main drivers underlying species richness patterns at large scales. However, as many landscapes and thus species' habitats are strongly influenced by humans, particularly in Central Europe, the effect of human land use on richness patterns cannot be neglected. The severe anthropogenic changes in land cover, increasing landscape fragmentation, degradation and intensification have already led to large losses in (semi-)natural vegetation, biodiversity and ecosystem functions and are expected to cause even more losses in the future.

Protected areas play a major role in the alleviation of anthropogenic pressures on species and their habitats. However, as site selection for many of these areas has not been based on knowledge on species occurrences, their effectiveness might be limited.

The present thesis, conducted in the federal states of Lower Saxony and Bremen in northern Germany, therefore aims at 1.) identifying the richness patterns of vascular plant species in the two states; 2.) testing the effectiveness of the protected area network for the conservation of threatened species in the lowland region of the study area and 3.) analysing the relationship between species richness patterns and their underlying processes with special regard to the human influence.

In order to address these aims, analyses were carried out on grid-based floristic data obtained from the vascular plant survey of Lower Saxony (*Niedersächsisches Pflanzenartenerfassungsprogramm*, NLWKN 1982-2003) that were combined with high-resolution data on land cover and information on environmental conditions and the location and extent of protected areas. In doing so, the thesis represents the first comprehensive analysis of the extensive floristic data base of Lower Saxony.

To analyse the patterns of vascular plant species richness across the study region, the overall species richness as well as the richness of five sub-groups of species aggregated using information on residence and threat status of the species inside each landscape unit (1,762 grid cells of c. 30 km²) was assessed. Richness of all species groups was found to be heterogeneously distributed across the region showing an increase in richness from north to south and – to some extent – also from west to east. The centres of high plant species richness of the different groups largely correlated with each other but also showed some interesting divergences.

The effectiveness of the protected area network in the lowland region of the state was shown to be generally high: 'Hotspots' of threatened species were found to be largely covered by nature reserves (German: *Naturschutzgebiete*) and Natura 2000 sites with the latter areas expediently complementing the former. It remained unclear, however, if the effective 'territorial' protection is reflected in an effective species protection.

The anthropogenic influence on the landscape of the study region was assessed using two separate analyses. These two studies focussed on the relationship between species richness and the degree of landscape fragmentation on one hand and landscape modification on the other hand.

The degree of landscape fragmentation was assessed by applying the *Effective Mesh Size Index* (m_{eff}). The relationship between m_{eff} and species richness of the total number of species as well as five sub-groups of vascular plants aggregated using their residence and threat status was assessed using a variation partitioning approach that separated the effects of landscape fragmentation, environmental conditions and spatial autocorrelation. Landscape fragmentation was found to significantly affect richness of all species groups. However, the strength of the effect varied between species, being largest for neophytes and smallest for threatened species.

A clustering analysis on landscape metrics describing landscape composition and configuration revealed a six-level gradient of landscape modification for the study region. The gradient spanned from highly fragmented urban landscapes to little fragmented landscapes with a high proportion of forest. Species richness of the total number of species as well as seven sub-groups of vascular plants aggregated using the residence and threat status as well as habitat preferences of the species were found to significantly differ along the gradient. For the entire and native species pool as well as for forest, low-nutrient indicating and threatened plants, species numbers peaked at the opposite ends of the modification gradient. For neophytes, urban plants and high-nutrient indicators richness decreased along the gradient from high species numbers in urbanised to low numbers in less modified landscapes.

The results obtained from the investigations in this thesis provide a basis for future landscape and conservation planning approaches in the study region by facilitating the identification of landscapes of high conservation value and/or concern. Provided that appropriate, high quality data like those used in the present thesis are available, the methodologies applied and developed in context of the thesis can be transferred to other regions to accordingly support conservation and landscape planning.

Zusammenfassung

Die auf verschiedenen Skalen-Ebenen heterogene Verteilung des Artenreichtums fasziniert Naturforscher und Biologen seit Jahrhunderten, so dass eine Vielzahl von Hypothesen zur Erklärung dieser Muster aufgestellt wurde. Es besteht breiter Konsens darüber, dass eine Vielzahl verschiedener Prozesse für die Variation des Artenreichtums verantwortlich ist. Je nachdem welche Taxa, welche Orte und welche räumlichen und zeitlichen Skalen betrachtet werden, wird den verschiedenen Faktoren dabei eine unterschiedlich große Bedeutung zugemessen. Umweltgradienten gelten in diesem Zusammenhang als die wichtigsten, den Artenreichtum bestimmende Faktoren. Da jedoch die Landschaften und damit Habitate der Arten weltweit und vor allem in Mitteleuropa stark anthropogen überprägt sind, sollte der auf den Artenreichtum wirkende Einfluss anthropogener Interventionen in diesem Zusammenhang nicht vernachlässigt werden. Die durch den Menschen und seine Aktivitäten verursachten Landnutzungsänderungen, eine steigende Landschaftsfragmentierung und -degradation sowie die Intensivierung der Landnutzung haben sich bereits im Verlust (halb-)natürlicher Landschaftselemente, dem Rückgang der Biodiversität und der Verschlechterung von Ökosystemfunktionen niedergeschlagen; eine weitere Verschlechterung der Zustände wird vorhergesagt.

In diesem Zusammenhang dient die Ausweisung von Schutzgebieten vielfach dazu, den auf die Arten und ihre Habitate wirkenden Druck zu vermindern. Da die Lage dieser Gebiete jedoch häufig nicht auf Grundlage der Kenntnis über das Vorkommen von Arten bestimmt wurde, dürfte die Effektivität der Schutzgebiete für den Artenschutz vielfach eingeschränkt sein.

Die vorliegende, die norddeutschen Bundesländer Niedersachsen und Bremen umfassende Arbeit untersucht 1.) die Muster des Artenreichtums der Gefäßpflanzenarten in den beiden Ländern; 2.) die Effektivität des Schutzgebietsnetzwerks des norddeutschen Tieflands für den Schutz seltener und bedrohter Gefäßpflanzenarten und 3.) den Zusammenhang zwischen Mustern des Artenreichtums und den ihnen zugrundeliegenden Prozessen unter besonderer Berücksichtigung des anthropogenen Einflusses. Die Arbeit basiert auf einem umfangreichen, im Rahmen des Niedersächsischen Pflanzenartenerfassungsprogramms (NLWKN 1982-2003) erhobenen Datensatzes, der räumlich explizite Informationen zur Verbreitung aller in den Bundesländern vorkommenden Arten liefert. Die floristischen Daten wurden für die durchgeführten Analysen mit hochaufgelösten Daten zu Landbedeckung und Umweltbedingungen kombiniert. Damit repräsentiert die vorliegende Arbeit die erste umfassende Auswertung des umfangreichen floristischen Datensatzes.

Für die Untersuchung der Muster des Gefäßpflanzenreichtums innerhalb der Untersuchungsregion wurden für die Gesamtzahl der Arten sowie für fünf auf Grundla-

ge des floristischen Status' und des Gefährdungsgrades der Arten zusammengestellte Untergruppen der Artenreichtum pro Landschaftseinheit (1.762 Messtischblatt-Quadranten à ca. 30 km²) berechnet. Der Artenreichtum aller Gruppen zeigte eine heterogene Verteilung in der Untersuchungsregion, wobei eine Zunahme der Artenzahlen von Nord nach Süd und zum Teil auch von West nach Ost erkennbar war. Die Zentren hoher Artenvielfalt der verschiedenen Gruppen korrelierten miteinander.

Die Analyse der Effektivität der Schutzgebiete für den Schutz der seltenen und bedrohten Gefäßpflanzenarten zeigte, dass die Artvorkommen im niedersächsischen Tiefland zu einem relativ hohen Anteil durch Naturschutz- und Fauna-Flora-Habitat-Gebiete abgedeckt sind, wobei letztere Gebiete die erstgenannten sinnvoll ergänzten. Im Rahmen der Analyse konnte jedoch nicht untersucht werden, inwiefern sich der nachgewiesene effektive Gebietsschutz in einem effektiven Artenschutz niederschlägt.

Der anthropogene Einfluss auf die Landschaft der Untersuchungsregion wurde im Rahmen zweier separater Analysen untersucht, wobei der Zusammenhang zwischen Artenreichtum und Landschaftsfragmentierung einerseits und dem Grad der Landschaftsmodifikation andererseits analysiert wurde.

Die Landschaftsfragmentierung wurde in der vorliegenden Arbeit mittels des Landschaftsmaßes der 'Effektiven Maschenweite' (m_{eff} ; engl. *Effective Mesh Size Index*) untersucht. Der Zusammenhang zwischen m_{eff} und dem Artenreichtum aller Gefäßpflanzenarten sowie fünf weiteren nach ihrem floristischen bzw. Gefährdungsstatus definierten Gruppen wurde mittels einer Varianzpartitionierung ermittelt, die die Effekte der Landschaftsfragmentierung von den durch Umweltvariablen und räumliche Autokorrelation der Daten verursachten Effekten separiert. Der Grad der Landschaftsfragmentierung erklärte einen signifikanten, jedoch unterschiedlich großen Anteil des Artenreichtums aller untersuchten Gruppen, wobei der stärkste Effekt für die Gruppe der Neophyten und der geringste für die Gruppe der seltenen und bedrohten Arten erkennbar waren.

Eine Cluster-Analyse auf Grundlage verschiedener, die Komposition und Konfiguration der Landschaft beschreibender Landschaftsmaße identifizierte für die Modellregion einen sechsstufigen 'Landschaftsmodifikations'-Gradienten. Dieser Gradient reichte von stark fragmentierten urbanen hin zu wenig fragmentierten Landschaften mit hohem Waldanteil. Der Gesamtartenreichtum sowie die Artenzahlen von sieben verschiedenen, nach ihrem floristischen Status, ihrer Gefährdung bzw. ihrer Habitat-Bindung bestimmten Artengruppen zeigten signifikante Unterschiede entlang des Gradienten. Die Gesamtartenzahl wie auch der Reichtum der indigenen, der bedrohten sowie der an Waldlebensräume und nährstoffarme Habitate gebundenen Arten war jeweils an den Enden des Gradienten am höchsten. Der Reichtum der Neophyten sowie der an urbane Räume und nährstoffreiche Habitate gebundenen Arten nahm dagegen von den urbanen zu den wenig beeinflussten Landschaften hin ab.

Die im Rahmen der Arbeit erhaltenen Ergebnisse können zukünftig als Grundlage für naturschutzfachliche Planungen in der Untersuchungsregion dienen, indem sie

die Ausweisung von für den Naturschutz relevante Landschaftsräume unterstützen. Sofern entsprechende, qualitativ mit denen der vorliegenden Arbeit vergleichbare Grundlagendaten vorliegen, eignen sich die im Rahmen der Arbeit angewendeten und entwickelten Methoden, um auf andere Regionen übertragen zu werden und auch dort naturschutzfachliche und landschaftsplanerische Prozesse zu unterstützen.

1

Introduction

1.1 Patterns of species richness at the landscape scale

The heterogeneous geographic distribution of species richness has fascinated biologists for centuries (RICKLEFS, 2004; see BARTHLOTT et al., 2005 for an overview of the history of the exploration of diversity patterns of vascular plants). The acceleration of extinction rates in recent decades (HODAPP et al., 2014) has meant that the identification and analysis of the spatial patterns of species richness is now more important than ever. Species richness is frequently referred to as an indicator for biodiversity (HODAPP et al., 2014) as it is comparatively easy to measure and does not depend on knowledge of the abundance of species. Biodiversity, in turn, is widely accepted as to play an important role for ecosystem functioning (SECRETARIAT OF THE CBD, 2006; CARDINALE et al., 2012; HODAPP et al., 2014). Therefore, species diversity has been used as a proxy for the ecological status of ecosystems (HODAPP et al., 2014).

The importance of biodiversity is widely acknowledged in science as well as in political decision making and therefore numerous measures to explore and protect it have been implemented. On the global scale, the Convention on Biological Diversity (CONFERENCE OF THE PARTIES, 1992) is ratified by 194 countries (status September 2014; SECRETARIAT OF THE CBD, 2014) and aims at the "conservation of biological diversity, the sustainable use of its components and the fair and equitable sharing of the benefits arising out of the utilization of genetic resources" (SECRETARIAT OF THE CBD, 2005, p. 5). These goals have been implemented in European legislation by the EU Biodiversity Strategy (EUROPEAN COMMISSION, 1998). In this context, the Habitats Directive (together with the Birds Directive) forms the "cornerstone of Europe's nature conservation policy" (EUROPEAN COMMISSION, 2014) and aims at the protection of biodiversity, i. e. the diversity in flora, fauna and habitats.

As only what is known can be effectively protected, the identification of species richness patterns is an important aspect of their effective protection. Plants are seen as the "key structural elements of terrestrial ecosystems" (KREFT & JETZ, 2007, p. 5925) as they play an important functional role as primary producers and provide habitats for animal species (CARDINALE et al., 2011; SCHETTER et al., 2013). Because of this, as well as the fact that data on plant species occurrences are relatively easy to collect compared to the collection of data of mobile organisms (SCHETTER et al., 2013), a lot of research has focused on these organisms. Patterns of vascular plant species richness have been investigated on the global (MYERS et al., 2000; BARTHLOTT et al., 2005; KIER et al., 2005) as well as on the European scale (ESSL et al., 2013) and have been used for global-scale conservation priority setting (MYERS et al., 2000).

Species richness patterns may be either directly derived by summarising the knowledge on the spatial distribution of single species (see e. g. HAEUPLER, 2000; GARVE, 2007) or – if reliable data are missing – estimated by applying modelling approaches (cp. KIER et al., 2005; MAES et al., 2005; KREFT & JETZ, 2007).

The availability of digital data on species occurrences on a regional and national level has increased rapidly in recent years. At the global scale, the Global Biodiver-

sity Information Facility (GBIF) provides access to more than 400 million spatially explicit records of several taxonomic groups, including vascular plants (GBIF, 2014). In Europe, most countries maintain species mapping projects that gather data on the distribution and presence or even abundance of plant and animal species (see e. g. ZAJAC & ZAJAC, 2001; LANDUYT et al., 2012; NETPHYD & BfN, 2013). The mapping of species occurrences in Germany is supervised by the federal states (see e. g. HARDTKE & IHL, 2000; KORSCH et al., 2002; HAEUPLER et al., 2003). The results of this mapping are then reported to the federal authority.

Data on plant species occurrences provide a good basis for conservation planning (BROOKS et al., 2004). Therefore information obtained from floristic mapping have been applied for various purposes. For example, HAEUPLER & VOGEL (1999) used data obtained from such a mapping to derive patterns of species richness for the whole of Germany. Based on species number of rare and threatened plants per landscape unit, the authors determined eight 'hotspots'¹ of phytodiversity distributed across the country. On a smaller scale, WINTER et al. (2010b) used information on the occurrences of threatened plant species in an administrative district of Germany to identify Important Plant Areas (IPAs) that form an integral part of the Global Strategy for Plant Conservation (THE SECRETARIAT OF THE CBD, 2002).

Species records have furthermore been used to analyse the indicative power of selected sub-groups of plant species for the overall species richness of a region: For north-west Germany, DIEKMANN et al. (2008) used records of forest species taken from a flora atlas of the Weser-Elbe region and related the richness of these species to that of the overall number of plants reported for the region. The authors showed that species numbers of this group are indicative for total species richness as well as for the number of threatened species. Similarly, KATI et al. (2004) found that woody plants acted as suitable indicators for the richness of other taxonomic groups in the Mediterranean.

In the study area of the present thesis, the federal states of Lower Saxony and Bremen, species records amongst other things serve as a basis for the creation of distribution maps², the compilation of Red Lists and as a contribution to landscape and environmental planning (SCHUPP et al., 2001).

1.2 Linking patterns and processes: What influences species richness patterns at the landscape scale?

The analysis of the underlying causes of species richness patterns is a major goal in ecological, evolutionary, and biogeographical research (GASTON, 2000; GODFRAY &

¹In the following the term 'hotspot' is used for landscape units hosting a high richness of threatened plant species (cp. definition of HAEUPLER & VOGEL, 1999).

²See GARVE (1994) for such maps for threatened plant species and GARVE (2007) for all vascular plant species of the state.

LAWTON, 2001; GOTELLI et al., 2009; WIENS, 2011) and has challenged researchers for centuries (RICKLEFS, 2004). In recent decades, researchers have provided many different ideas on the question of what influences richness patterns (see RICKLEFS, 2004 for an overview on the history of these concepts). In accordance with many researchers, COLWELL et al. (2004, p. E14) suggests that "multiple factors are surely at work and that any particular factor may be more or less important for different taxa, different places, and different spatial and temporal scales". Processes and factors influencing species richness patterns include speciation and extinction rates (COLWELL et al., 2004; WIENS, 2011), geometric constraints (i. e. mid-domain effect, MDE, COLWELL & HURTT, 1994) as well as geographic patterns of temperature, productivity, potential evapotranspiration, topography, formation history (COLWELL et al., 2004) and human activities (WHITE & KERR, 2007; WANG et al., 2011). The following two subsections will focus on the role of abiotic environmental conditions and the human influence shaping species richness patterns. It has to be kept in mind, though, that the underlying processes influencing these patterns are far too complex to be fully disentangled. Thus, usually the processes are approached by using indicators or surrogates.

1.2.1 Environmental conditions as key drivers for richness patterns of plants

The role of the abiotic environment in shaping species richness patterns has been analysed on various spatial scales ranging from global (KREFT & JETZ, 2007) to local assessments (HEIKKINEN, 1996). BARTHLOTT et al. (2007, p. 308) describes 'geodiversity', i. e. "the heterogeneity of topography, geology, soils, or climate", as the main driver for global spatial patterns in species richness. More specifically, KREFT & JETZ (2007) showed that for vascular plants potential evapotranspiration, the number of wet days per year, and measurements of topographical and habitat heterogeneity are core predictors of global richness patterns. Similar factors were found to explain species richness on a smaller scale: Elevation range followed by minimum monthly potential evapotranspiration was identified as to be the most important factor determining woody plant species richness of the Iberian Peninsula VETAAS & FERRER-CASTÁN (2008). Similarly, climatic variables like temperature and potential evapotranspiration were found to be by far the most important predictors for vascular plant species richness in Austria (MOSER et al., 2005). For alien plant species richness in the European Alps, environmental heterogeneity proved to be the best surrogate (DAINESE et al., 2014). The same factor in combination with the presence of calcareous rock outcrops and threshold values of minimum precipitation was found to also determine species richness in Switzerland (WOHLGEMUTH, 1998). For Germany, native and alien plant species richness was found to be largely dependent on the diversity of geological substrates (KÜHN et al., 2003). Similarly, on the local scale the overall habitat diversity was identified as the main driver for species richness of vascular plants in a nature reserve in northern Finland (HEIKKINEN, 1996).

All these findings promote the hypothesis that overall species richness tends to be higher in a varied environment compared to a monotonous one, as heterogeneous environmental conditions provide a greater diversity of niches potentially suitable for plant species (HONNAY et al., 2003; ADAMS, 2009).

1.2.2 The human impact on richness patterns of plants

The previous subsection dealt with the role of the abiotic environment in shaping species richness patterns. Undoubtedly, environmental conditions are the main drivers underlying species richness patterns on a large scale. However, as many landscapes and thus species' habitats worldwide and particularly in Central Europe are strongly influenced by humans (KOWARIK, 1990), the shaping effect of human interventions on richness patterns should not be overlooked.

Over the last few decades, much attention has been focused on the analysis of the effects of human interventions at the landscape scale. In this context, a large number of studies have searched for surrogates describing the human impact on landscapes and biodiversity.

Human-induced threats to and protection of plant species richness

Habitat loss, fragmentation and degradation have been identified as being the main factors threatening species richness (FAHRIG, 2001; AGUILAR et al., 2006; KRAUSS et al., 2010). Both habitat fragmentation and loss – the latter being an "important by-product of habitat fragmentation" (MCGARIGAL, 2014, p. 89) – are known to increase the isolation between plant or animal sub-populations, thereby impeding the genetic exchange between individuals. The resulting loss in genetic variation may increase the probability of extinction by demographic and environmental stochasticity (BOOY et al., 2000). In many cases, habitats are not immediately fully destroyed but undergo a gradual degradation that, however, ultimately leads to their complete destruction.

A multitude of landscape-level processes are responsible for habitat loss, fragmentation and degradation in Western Europe, including – but not being limited to – urbanisation, agricultural intensification, eutrophication and climate change (cp. Fig. 1.1).

Urbanisation³ has been identified as one of the leading causes of species extinction (MCKINNEY, 2006; ARONSON et al., 2014). Urbanisation affects biodiversity in many ways, e. g. through the fragmentation and destruction or degradation of species' habitats and by altering the precipitation and temperature regime as well as air, soil and water quality (SUKOPP & STARFINGER, 1999). In their review on causes of species endangerment in the United States, CZECH et al. (2000) found that urbanisation in the country endangers more species than any other human activity although barely 5%

³i. e. "the process by which towns and cities are formed and become larger as more and more people begin living and working in central areas" (Definition taken from MERRIAM-WEBSTER (2014)).

of the total surface area of the United States is covered by urban and other built-up areas (MCKINNEY, 2002). In Germany, urban and traffic areas cover about 14 % of the landscape (STATISTISCHE ÄMTER DES BUNDES UND DER LÄNDER, 2012), and this proportion is increasing: Each day about 80 ha of undeveloped land are transferred to building land (UBA, 2014) resulting in an enormous loss of unsealed surface area and a fragmentation and loss in species' habitats. Despite the destructive effect of urbanisation on native habitats, cities are found to be generally richer in plant species than surrounding areas (HONNAY et al., 2003; WANIA et al., 2006; LIPPE & KOWARIK, 2008). This pattern might arise from different aspects: Generally, niche availability in urban areas is known to be rather high (DEUTSCHEWITZ et al., 2003; HONNAY et al., 2003; WANIA et al., 2006). KÜHN et al. (2004) furthermore suggested that urban areas are commonly situated in naturally diverse ecoregions resulting in an overall increased species diversity. Also, alien species are known to show increased numbers in urban areas, adding to the urban species pool (ROY et al., 1999; KÜHN et al., 2004).

Agricultural intensification has been proposed as one further main cause of biodiversity loss (TILMAN et al., 2001; TSCHARNTKE et al., 2005). In Central Europe it accelerated in course of industrialisation in the late 19th and early 20th century (TSCHARNTKE et al., 2005; KRAUSE & CULMSEE, 2013) and has been reinforced by the advent of the common agricultural policy of the EU since the 1950s (STOATE et al., 2001; KRAUSE et al., 2011). Conventional farming practices affect species diversity mainly through the conversion of non-arable land that increases habitat loss and fragmentation (TSCHARNTKE et al., 2005). However, not only the conversion of non-arable to high-productive cropland but also the abandonment of marginal sites, the removal of small-scale landscape elements and the application of pesticides and fertilisers have an effect on species (TILMAN et al., 2001). Studies on the landscape scale in Europe have identified large losses in plants, invertebrates and farmland birds following agricultural intensification (ANDREASEN et al., 1996; DONALD, 1998; DONALD et al., 2001; BUTLER et al., 2010; STORKEY et al., 2012; MEYER et al., 2013).

The **eutrophication** of ecosystems, i. e. primarily the input of nitrogen, phosphorous and sulphur compounds, has been identified to reduce plant species richness in a broad range of European ecosystems (DISE, 2011). As high fertiliser inputs largely add to the eutrophication of a landscape, the process is tightly linked to that of agricultural intensification. Additional inputs adding to the eutrophication of ecosystems come from industry and transport (UNEP, 2014). Due to strict regulations in Europe (see 'Convention on Long-range Transboundary Air Pollution' and its eight subsequent protocols; UNECE, 1979), emissions of air pollutants, especially sulphur, have been reduced in recent years (DISE, 2011). However, critical loads are still exceeded in many semi-natural ecosystems in Europe (HETTELINGH et al., 2008; DISE, 2011). Eutrophication affects species mainly through the degradation of their habitats – in the worst case habitats will become uninhabitable for the species under consideration which results in habitat loss. The negative impacts of eutrophication on species

richness have been reported for all kinds of habitats across Europe, e. g. for acid grasslands in the Atlantic biogeographic region (STEVENS et al., 2010), southern Swedish temperate oak forests (BRUNET et al., 1998), Swiss mountain grasslands (ROTH et al., 2013) and British heathlands as well as acid and calcareous grasslands (MASKELL et al., 2010).

Based on different scenarios, the impact of **climate change** on the distributions of plant and animal species has been frequently modelled in recent years (e. g. BAKKENES et al., 2002; THUILLER et al., 2005; POMPE et al., 2008). These analyses have shown that many European plant species are at risk from climate change (THUILLER et al., 2005). THOMAS et al. (2004) predicted that due to climate change 6–8 % of all plant species in Europe could become extinct by 2050. For Germany, POMPE et al. (2008) suggested that – even under moderate climate change assumptions – negative impacts on the German flora will arise. The authors point out that due to climate change severe shifts in the species pools will occur.

Climate change scenarios up to the year 2080 for Germany have identified a definite warming trend of 1.6–3.8 °C and small overall changes in annual precipitation (<10 %) (ZEBISCH et al., 2005). However, changes in seasonal precipitation will occur with an increase in winter and a decrease in summer precipitation (ZEBISCH et al., 2005). Due to their particular environmental characteristics, different regions in Germany are assumed to be affected by climate change by a varying degree (ZEBISCH et al., 2005). Highest vulnerability to climate change is expected for south-west and the central parts of eastern Germany as well as the Alps. The low mountain ranges of Germany as well as the coastal regions show medium vulnerability whereas north-west Germany was identified to be least vulnerable (ZEBISCH et al., 2005). Therefore, impacts of climate change on the study area of this thesis, the federal states of Lower Saxony and Bremen, can be assumed to be small compared to other German states.

The processes discussed in the previous paragraphs all add to the modification of a landscape and by doing so pose a threat to species. As it is difficult, if not impossible, to consider all of these processes separately as their effects on biodiversity in a landscape are usually interlinked, multi-layered concepts combining them have been proposed by several researchers. These concepts aim at describing the degree of modification a landscape has undergone. Concepts proposed involve 'landscape modification' (FORMAN & GODRON, 1986) and the related concepts of 'landscape naturalness' (MACHADO, 2004; LIIRA & SEPP, 2009; WINTER et al., 2010a), 'hemeroby' (SUKOPP, 1976; KOWARIK, 1988; GRABHERR et al., 1998; STEINHARDT et al., 1999; WRBKA et al., 2004; WALZ & STEIN, 2014), 'urbanity' (HILL et al., 2002; WRBKA et al., 2004) and the 'degree of culturalness' (JANSEN et al., 2009). These concepts are used to assess the conservation value of landscapes (WRBKA et al., 2004) and have been proposed to allow conclusions on biodiversity (WALZ & STEIN, 2014). However, until now no study has directly analysed the relationship between any of these concepts and species richness/diversity.

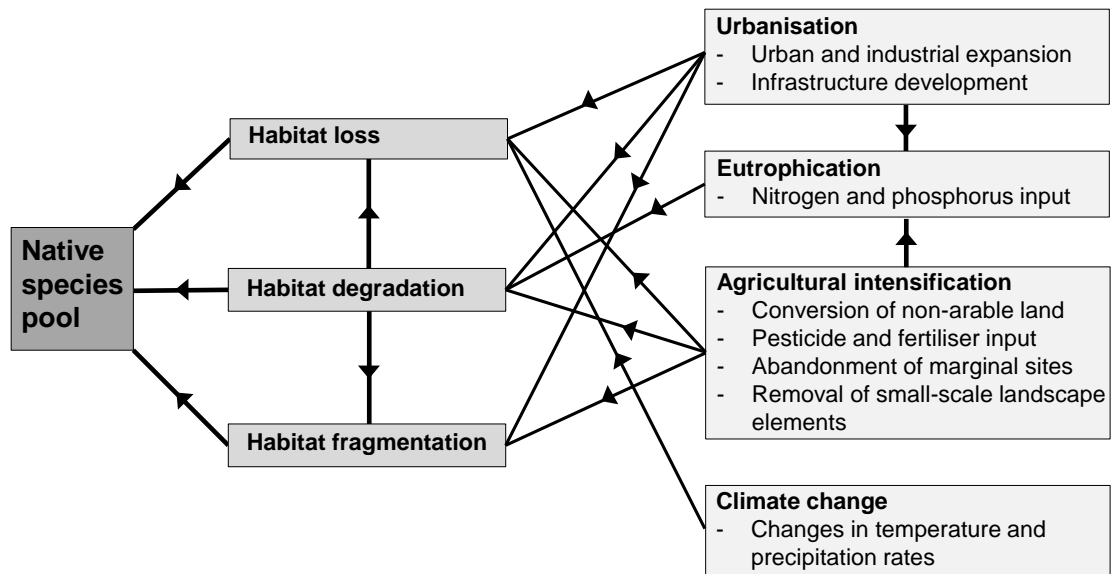


Figure 1.1: Human-induced threats to the local species pool (selection of main threats). Arrows indicate the main direction of the impact.

As discussed in the previous paragraphs, species in today's highly industrialised landscapes are threatened by a multitude of processes, most of them being linked to human-induced changes in the landscape. To reduce the negative impacts of these changes on species, a number of different concepts for the protection of species have been proposed that can generally be allocated into segregative and integrative measures (PLACHTER, 1991). These concepts have been developed for agricultural landscapes but can analogously be transferred to forests, water bodies and urban areas (PLACHTER, 1991). In segregative concepts, areas used for production are separated from those designated for the protection of species, habitats and ecosystems (JESSEL, 2012). In contrast, integrative concepts try to include both production and conservation in the same areas. Integrative measures reduce negative impacts on species by a reduction in the intensity of land use and the establishment of biotope networks (PLACHTER, 1991). This management can, amongst others, promote species tolerant to land use (TSCHARNTKE et al., 2012), improve ecosystem services and reduce isolation of habitats (JESSEL, 2012; see also "dual conservation strategy", JACKSON et al., 2009). However, it can not promote the protection of species linked to (semi-)natural ecosystems (e. g. mires, rocky outcrops). For the protection of such species the segregative concept might be useful. Segregation is mainly enforced by the designation of protected areas inside which land use is forbidden, restricted or regulated to provide habitats for species otherwise lost from a region.

In Germany, there are various categories of protected areas, of which national parks, nature reserves (German: *Naturschutzgebiete*) and Natura 2000 sites (EU Habitats Directive) are those covering the largest areas. Protected areas are widely ac-

cepted as being essential for the long-term conservation of biodiversity (EVANS et al., 2006). However, their effectiveness in protecting species remains largely unclear as the extent and location of many protected areas have not been defined using knowledge on species occurrences (WINTER et al., 2010b; JEDICKE, 2012). Determining the effectiveness of such areas "in representing and maintaining biodiversity is a core issue in systematic conservation planning" (GASTON et al., 2006, p. 77). Nevertheless, few studies until now have focused on this topic. Regional studies that analysed the effectiveness of protected areas have identified numerous gaps in the existing protected area network (see e.g. SCOTT et al., 2001 for the USA, ANDELMAN & WILLIG, 2003 for the Western Hemisphere and DEGUISE & KERR, 2006 for Canada). An analysis by RODRIGUES et al. (2004) uncovered such gaps on a global scale. In contrast, for Great Britain, JACKSON et al. (2009) found the current protected areas to cover the occurrences of a large majority of Red List plant species. However, despite the good overall species coverage, less than one third of the total number of occurrence records were covered. It is obvious that results obtained for one particular region cannot be directly transferred to other areas and that further research in this field is strongly needed.

Surrogates for the human influence on species richness patterns

As mentioned earlier, much attention has focused on the analysis of the effects of human interventions on species richness patterns in recent decades. Studies on this topic usually have searched for surrogates describing and quantifying the human impact on landscapes and biodiversity. For this purpose, land cover data that reflect the influence of humans on the landscape as well as landscape metrics used to quantify the magnitude of this impact have been widely applied. Using landscape metrics either the composition (i. e. the abundance of patch types in a landscape) or configuration (i. e. the spatial arrangement of patches) of a landscape can be assessed (cp. subsection 2.2.5). Indices used include measures of landscape diversity, patch shape complexity, edge length and density, core area sizes as well as metrics used to quantify landscape fragmentation, landscape/habitat connectivity and isolation. Below, some of these measures and their relevance for species richness and diversity are discussed.

Landscape diversity – analogical to 'geodiversity' (cp. subsection 1.2.1) – was identified to be a convenient surrogate for species richness (LUOTO, 2000; HONNAY et al., 2003). Different measures have been used to express landscape diversity, the simplest merely reflecting the number (i. e. patch richness) or proportions of different land cover types (DEUTSCHEWITZ et al., 2003; KÜHN et al., 2003; WANIA et al., 2006), the more complex ones combining richness and evenness of land cover types (e. g. Simpson's and Shannon's Diversity Index; see e. g. DEUTSCHEWITZ et al., 2003; HONNAY et al., 2003; KUMAR et al., 2006).

Several studies have analysed the relationship between species richness and landscape diversity. For Germany, a high structural and habitat diversity in combination with moderate levels of natural and/or anthropogenic disturbance was found to increase richness of native and alien plant species (DEUTSCHEWITZ et al., 2003). Similarly, regional plant species richness of Flanders, Belgium was found to be positively correlated with landscape diversity expressed in terms of high patch richness and evenness and irregular patch shapes (HONNAY et al., 2003). As landscape diversity in the study area was found to go along with the degree of urbanisation also a positive correlation between the percentage of built-up area and species richness was stated. As previously discussed in the paragraph on the effects of urbanisation, the latter has been frequently reported by researchers and has been attributed to the increased niche availability in urban areas.

However, not only landscape diversity measures have been applied to explain species richness patterns. Several studies have used **patch shape complexity** as a surrogate for species richness as they assumed that simple patch shapes will be dominant in landscapes strongly influenced by humans, whereas more natural landscapes should be characterised by irregular shaped patches (MOSER et al., 2002; YAMAURA et al., 2008). In the study of MOSER et al. (2002) an index of patch shape complexity developed by the authors that is based on the number of shape-characterising points along a polygon's boundary proved to be a good predictor for the species richness of vascular plants and bryophytes in Austria: the authors found species richness to be highest in landscapes for which irregular patch shapes were reported. Similarly, for a Mediterranean forest landscape SCHINDLER et al. (2013) identified patch shape to be a good indicator for the diversity of woody plants. In contrast, by using a shape index independent of area for Japan, the study of YAMAURA et al. (2008) found the richness of forest-floor plants to be influenced by patch shape to only a limited extent whereas local forest structure had a much larger effect.

As **landscape fragmentation** resulting in habitat loss and isolation is supposed to be one of the largest threats to biodiversity worldwide (JAEGER, 2000; see also previous section on threats to species) its indicative power for species richness has been analysed by numerous studies at the local and landscape scale (see for an overview DEBINSKI & HOLT, 2000; FAHRIG, 2003). The simplest way in which the fragmentation of a landscape can be expressed is by using measures of patch numbers or patch density (MCGARIGAL, 2014). However, as these simple fragmentation measures are supposed to be insensitive and inconsistent in their behaviour (MCGARIGAL, 2014), attempts have been made to develop alternative and more explicit measures that "behave in a consistent and logical manner across all phases of the fragmentation process" (MCGARIGAL, 2014, p. 131). Still, despite their qualities, these indices (i. e. *Landscape Division Index*, *Splitting Index* and *Effective Mesh Size*; JAEGER, 2000) until now have not been applied as potential surrogates for species richness. Instead, frequently, the size and the **isolation** of patches or the **connectivity** between them were used as (inverse)

measure of fragmentation (e. g. HELM et al., 2006; COUSINS et al., 2007; KOLB, 2008; ZIMBRES et al., 2013). RAATIKAINEN et al. (2009) for extensive grasslands in southern Finland used a connectivity index taking into account the distances to all possible source populations ('IFM measure', cp. MOILANEN & NIEMINEN, 2002) to analyse how richness of spring ephemerals is related to habitat area and connectivity. They found the number of ephemerals to be highest in larger and well-connected study patches. In the study of ADRIAENS et al. (2006) a similar measure was applied to analyse how vascular plant species richness of calcareous grasslands in southern Belgium is influenced by landscape configuration. The authors found that the connectivity between grassland patches only affected the number of specialist species but not that of generalists. The opposite result was obtained by CRISTOFOLI et al. (2010) for heathlands in the high Ardenne (Belgium): By using measures of patch connectivity and isolation the authors found generalist, not specialist species richness to be correlated with connectivity. Interestingly, for islands in a large lake in the south-east of China, island isolation measured as the Euclidean distance to the mainland did not significantly explain vascular plant species richness (HU et al., 2011).

The aforementioned examples show that effects of landscape fragmentation and habitat isolation may vary depending on the species groups and habitats considered (EWERS & DIDHAM, 2006; RODRIGUEZ-LOINAZ et al., 2012). Therefore, further studies on this important topic using different measures of fragmentation and isolation/connectivity should be undertaken for different landscapes and habitats.

1.3 General objectives and thesis outline

The present thesis applies spatially explicit data on the distribution of vascular plant species in combination with various information on land cover and environmental conditions to identify processes shaping species richness patterns. In doing so, the analyses focus on the significance of the anthropogenic influence and only marginally consider the importance of environmental conditions. All analyses were conducted for the federal states of Lower Saxony and Bremen, northern Germany, for which spatially explicit area-wide data on the occurrences of all vascular plant species (plant survey of Lower Saxony, NLWKN 1993–2003) as well as information on land cover and environmental conditions are available.

The main goals of the thesis are:

- a) To identify and describe the patterns of vascular plant species richness in Lower Saxony and Bremen, Germany.
- b) To evaluate the adequacy of the protected area network for the protection of rare and threatened plant species in the lowland region of the study area.
- c) To analyse the relationship between richness patterns and the human influence on the landscape.

The thesis is subdivided into four main data chapters (chapters 3 to 6) briefly presented in the following paragraphs. These chapters are precluded by a chapter that introduces the reader to the study area and gives an overview on the data used in the analyses (chapter 2). The thesis closes with chapter 7 which summarises and synthesises the main findings of the previous chapters and highlights open questions and future research needs.

As the key conservation work and data collection is conducted by local authorities and the large number of volunteers working for the plant survey of Lower Saxony, considerable parts of the thesis (Chapter 3 and 4) were published in German to make the results accessible to these people in charge, of which not all are familiar with the English language. This audience-focused approach guarantees that the published research can be directly implemented within the study area. The German articles are precluded by detailed English summaries to make the key results available to the global research community.

Patterns of vascular plant species richness As pointed out in section 1.1, the identification of spatial patterns of species richness provides an important key aspect of conservation planning. The study presented in chapter 3 of this thesis makes use of the extensive species record database of the vascular plant survey of Lower Saxony and Bremen to derive richness patterns of six different species groups. The groups considered in the analysis comprise (a) the total number of species that has been reported for Lower Saxony and Bremen, (b) species grouped by their residence status (native species, archaeophytes, neophytes) and (c) species grouped by their threat status (non-threatened species, threatened species). The study serves as a starting point on which the studies presented in the following chapters are based.

The following research questions are answered by interpreting maps of the richness patterns of the aforementioned species groups and by taking into account the results obtained from a correlation analysis:

- Which patterns of vascular plant species richness are observable in Lower Saxony and Bremen and the three natural landscape regions, respectively?
- Do the different plant species (sub-)groups show richness patterns that are distinguishable from each other and how much do their richness centers overlap?
- In which areas do threatened species show highest species numbers and how do these hotspots differ from that of non-threatened species?

Adequacy of the protected area network As knowledge on the spatial distribution of the rare and threatened species of Lower Saxony and Bremen is of major importance for their effective protection, occurrences of these species have been recorded at a much finer resolution (c. 2 km²) than those of all other species. In the study presented in chapter 4 these high-resolution data were used to test to which extent the hotspots of rare and threatened species ('RL species') in the lowlands of Lower Sax-

ony are covered by protected areas. The analysis considers nature protection areas (*Naturschutzgebiete*, NSG) and NATURA 2000 sites. Additionally to the total number of threatened plant species Red List species linked to forest habitats ('RL forest species') are included in the analysis. The data are analysed using Generalised Linear Models (GLM) with Poisson distribution. The study aims at answering the following research questions:

- What patterns of diversity can be observed for the two species groups?
- Are occurrences of threatened species covered by nature protection areas?
- Does the designation of NATURA 2000 sites improve the situation?
- Are the results obtained for RL species transferable to RL forest species?

The human influence on species richness patterns As outlined in section 1.2, species richness patterns in industrialised countries are assumed to be strongly affected by human activities. To understand the magnitude and direction of this influence and the underlying regulatory processes is an important field of research. The two chapters dealing with this subject (chapter 5 and 6) use different methodological approaches to illuminate the relationship between species richness patterns and human influence.

The study presented in chapter 5 investigates whether landscape fragmentation expressed by means of the *Effective Mesh Size Index* is related to the richness of six different groups of vascular plants aggregated using information on their residence and threat status, respectively. The index has frequently been suggested to be ecologically meaningful (JAEGER, 2000; JAEGER et al., 2008; GIRVETZ et al., 2008) and is widely applied in studies on landscape fragmentation. However, its relevance for biodiversity has not been tested and its suitability as an indicator of biodiversity has even been questioned (ESSWEIN, 2007). The presented study aims to shed light on this topic by applying variation partitioning techniques used to disentangle the relationship between species richness, landscape fragmentation and environmental variables. Spatial autocorrelation in the data is considered by using PCNM analysis. It is hypothesised that:

- Landscape fragmentation, expressed as Effective Mesh Size (m_{eff}), varies in space, which results in a gradient in the degree of fragmentation within the sample of landscape units used in this study.
- The choice of fragmentation geometries that are used for the calculation of m_{eff} is of major importance for explaining plant species richness patterns on the landscape scale.
- Plant species richness varies between landscape units along the fragmentation gradient, with the effect size differing between different groups of species dependent on their residence (natives, archaeophytes and neophytes) and threat status (threatened vs. non-threatened).

The study presented in chapter 6 makes use of k-means clustering to derive a gradient of 'landscape modification' (cp. subsection 1.2.2) for 1386 landscape units of Lower Saxony and Bremen. The analysis is based on landscape metrics quantifying landscape composition and configuration. Clusters differing in terms of their degree of landscape modification are set up and placed along a 'gradient of landscape modification' according to the one proposed by FORMAN & GODRON (1986). The gradient is then linked to richness patterns of the total number of vascular plant species as well as that of seven sub-groups that were aggregated using information on residence and threat status and habitat preferences of the species. The study seeks to verify the following hypothesis:

- Based on landscape metrics used to assess the composition and configuration of a landscape, clusters of landscapes with similar characteristics can be differentiated. These can be placed along a gradient of landscape modification as proposed by FORMAN & GODRON (1986), spanning from (semi-)natural (mostly forested) to urbanised landscapes.
- The landscape clusters are meaningful for vascular plant species richness, i. e. species numbers vary significantly along the gradient of landscape modification intensity.
- Species numbers of different vascular plant species groups show different patterns along the gradient of landscape modification intensity.

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2 | Description of data

2.1 Study region

The study region comprises the German federal states of Lower Saxony and Bremen. Lower Saxony encompasses an area of 47.624 km² whereas Bremen as the smallest federal state of Germany has a size of only 404 km². As landscape formation history and natural environmental configuration, colonisation history and current land use vary widely across the state, Lower Saxony harbours a high number of different habitats. The area may roughly be subdivided into three regions, the coastal, the lowland and the upland area (Fig. 2.1), that largely differ in their characteristics.

The **coastal area** is dominated by marine deposits from the Holocene (HEUNISCH et al., 2007). It comprises the Wadden Sea, its islands and salt marshes, the estuaries of the three largest rivers of Lower Saxony (Ems, Weser and Elbe) as well as impoldered tidal wetlands. It shows an oceanic climate with mean annual precipitation rates of about 700–900 mm and mean annual temperatures of c. 9 °C. Main soil types occurring in this area are Fluvisols, Leptosols and Gleysols (NLFb, 1997¹). The coastal area is predominantly covered by grasslands with a proportion of 60 % whereas cropland areas contribute 20 % and forests merely 1 %.

The **lowland area** was formed during the Pleistocene and harbours various types of landscapes ranging from intensively managed agricultural landscapes in the western part to more wooded landscapes in the eastern part. From West to East, the climate changes from oceanic (annual precipitation rate c. 900 mm) to suboceanic-subcontinental (annual precipitation rate c. 550 mm). Mean annual temperatures in this area fluctuate around 8 °C. The climatic gradient is reflected in the soil characteristics: In the West the dominating soil types are those affected by seepage water, water logging and mass transfer (Histosols, Podzols); in the East primarily Cambisols and Leptosols are found. The area is dominated by croplands that cover about 40 % of this region; forests and grasslands are present at proportions of about 20 % each.

The **upland area** is a very heterogeneous region that is dominated by deposits from the Meso- and Palaeozoic (HEUNISCH et al., 2007). Due to its diverse topography, climatic conditions vary strongly. The uplands are dominated by aeolian silt as well as clay, sand and lime stone deposits (NLFb, 1997). Luvisols and Cambisols are the most common soil types in this region. Mean annual temperatures range between 8–9 °C. Mean annual precipitation rates vary between 650–750 mm in the lower parts of the area and 800–900 mm in the elevated parts. The Harz mountains at the South-western border of Lower Saxony feature very specific climatic conditions: The most elevated parts (up to 971 m a.s.l.) experience a montane climate with an annual mean precipitation rate of up to 1,500 mm and annual mean temperatures of only 7 °C (NLFb, 1997). The Harz region is dominated by base poor silicate bedrock as well as argillaceous shale and greywacke. Main soil types occurring in this area are Cambisols and Podzols (NLFb, 1997). Out of the three regions, the upland area features the high-

¹Translation of German soil types follows IUSS WORKING GROUP WRB (2006).

est proportion of forests (c. 30%); grasslands only prevail in about 9% of the area whereas croplands dominate c. 45% of the region.

The study region was subdivided into 1762 grid cells (*Messtischblatt-Quadranten*) with a size of about 5.5 km × 5.5 km (1/4 of a topographical map, scale 1 : 25,000; Fig. 2.1). Out of these 1386 grid cells are fully covered by land, 353 grid cells are located on the edge and 23 grid cells are fully covered by sea. For a more detailed study (see chapter 4, SCHMIEDEL et al., 2013), the lowland area of the state was overlaid with a more detailed grid of 15.729 cells with a size of about 1 km × 2 km (*Minutenfelder*, 1/60 of a topographical map, scale 1 : 25,000).

2.2 Data and data preparation

Data available for the study region included information on the distribution of vascular plant species, data on environmental variables as well as land-use types (Table 2.1). The following paragraphs give a detailed overview of these data.

Species data were analysed using MS Access (MICROSOFT CORPORATION, 2010) and ArcGIS-ArcInfo 9.2 (ESRI INC., 2007). ATKIS data, areal information as well as environmental variables – unless stated otherwise – were analysed using ArcGIS-ArcInfo 9.2, Hawth's Tools (BEYER, 2004) and Fragstats 3.3 (MCGARIGAL & CUSHMAN, 2002).

2.2.1 Species data

Occurrence data of all vascular plant species of Lower Saxony and Bremen were made available as an MS Access database by the vascular plant survey of Lower Saxony (*Niedersächsisches Pflanzenartenerfassungsprogramm*, NLWKN; Table 2.1). The data base (last update 15 December 2008) comprises 2,395,045 entries and covers the time span from 1993–2003 (first and second period of the state-wide mapping; third period is still underway). Out of all taxa listed in the data base only the 1879² non-erratic ones were considered³. Occurrence data were available on the grid basis mentioned in the previous chapter.

Prior to the analyses, species were aggregated on the highest taxonomical level found in the data base, i. e. subspecies were assigned to the species level and species to the respective aggregate found in the data base⁴. This aggregation affected 213 taxa (see Table A.3 in Appendix A for an overview). Thus, in total, 1666 taxa remained in the analysis out of which 72 were occurring in grid cells fully covered by sea or located on the edge of the study area only⁵ (cp. section 2.1) and thus were excluded

²Due to errors in the database that were detected just after the first article (SCHMIEDEL et al., 2011) was published, the total number of species considered in this article was 1819 taxa only.

³In total, 2708 taxa are listed in the floristic list of the data base out of which 829 are extinct, erratic or do not occur.

⁴The exception being the first article (SCHMIEDEL et al., 2011) which did not apply any aggregation.

⁵Hereafter referred to as 'edge only' species.

for the analyses conducted in SCHMIEDEL & CULMSEE (submitted) and SCHMIEDEL et al. (submitted).

For the analyses conducted in context of the present thesis, the species listed in the data base were aggregated into sub-groups using the residence and threat status and habitat preferences of the species, respectively (Table 2.2).

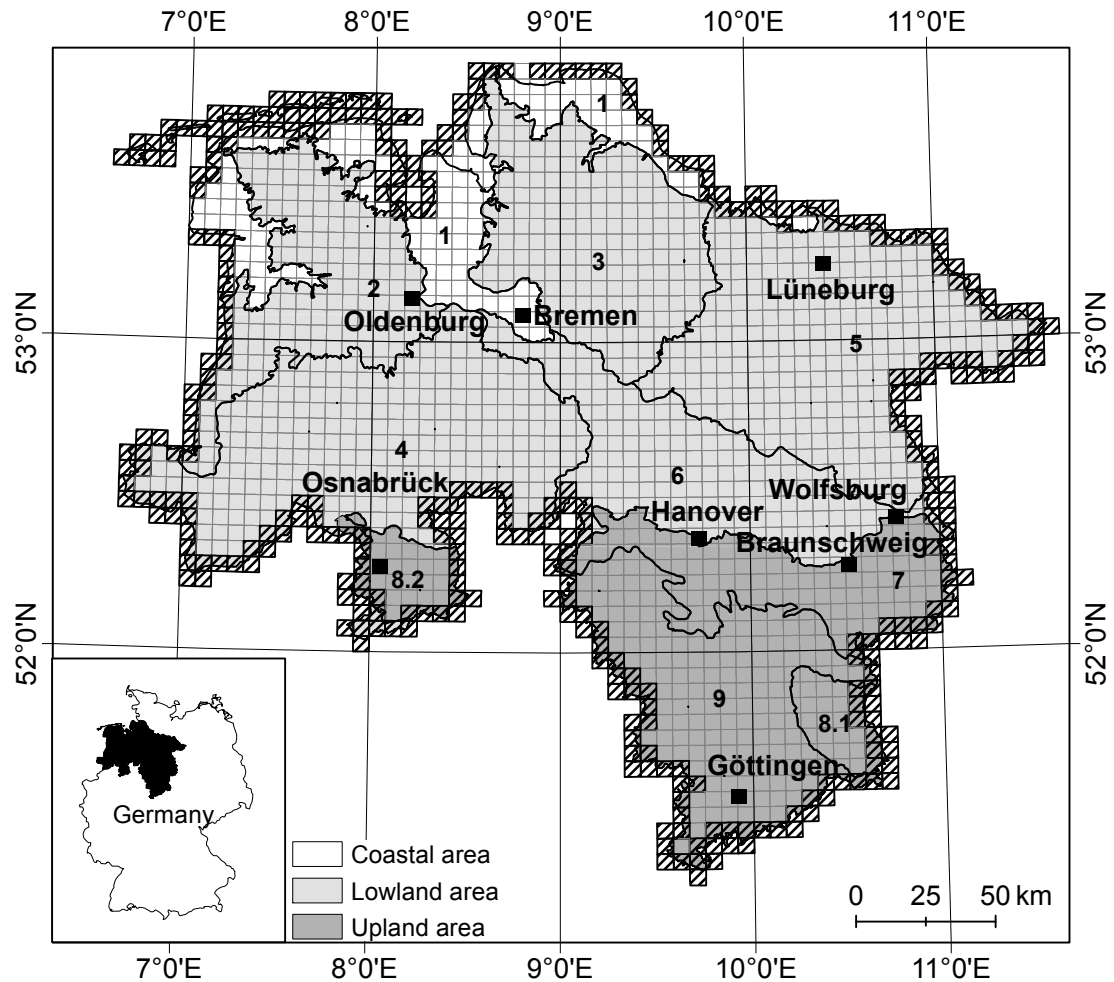


Figure 2.1: The study area encompassing the federal states of Lower Saxony and Bremen, with its three natural landscape regions (coastal, lowland and upland area). The region can be further subdivided into 10 bio-geographical regions (1 = Wadden Sea and marshes, 2 = East Frisian and Oldenburg moraine lowlands, 3 = Stade moraine lowlands, 4 = Ems, Hunte and Dümmer moraine lowlands, 5 = Lüneburg Heath and Wendland, 6 = Weser and Aller plains, 7 = Old moraine loess landscapes, 8.1 = Uplands of Osnabrück, 8.2 = Uplands of the Weser and Leine valleys, 9 = Harz mountains.) Plant occurrence data from the plant survey of Lower Saxony were available for 1762 quadrants of the topographical map (scale 1:25,000, i.e. about 5.5 km × 5.5 km in size). The area of hatched quadrants has only been partly inventoried, because they are either only partially covered by land area or are situated in neighbouring federal states.

The residence status of species was determined by using the information provided by the data base itself and information from KLOTZ et al. (2002): Native species were distinguished from archaeophytes (species that became naturalised prior to the 16th century) and neophytes (species that became naturalised in the 16th century or later) (see Table 2.2). Information on the threat status of species was derived from the Red List of Lower Saxony and Bremen (GARVE, 2004): Threatened species (Red List status 1, 2, 3, G and R) were distinguished from non-threatened species (Red List status V and *) (see Table 2.2).

Information on habitat preference, i. e. the linkage to urban and forest habitats were taken from KLOTZ & KÜHN (2002), BfN (2013) and SCHMIDT et al. (2011), respectively (Table 2.2). Urban species were defined as those species with an urbanity indicator value of 4 (urbanophilic) and 5 (moderately urbanophilic). Forest species are those species categorised as to occur in closed forests only (category 1.1, SCHMIDT et al., 2011) as well as forest edges and clearings (category 1.2, SCHMIDT et al., 2011). Information on Ellenberg Indicator Values (EIV) for nutrients (N) were taken from BfN (2013). Those species having a N value of 1 to 3 were defined as species of nutrient-poor habitats, those with a N value of 7 to 9 as species of nutrient-rich habitats (Table 2.2).

Table A.2 in Appendix A lists all species, along with their residence and threat status as well as their habitat preferences and furthermore denotes the 'edge only' species.

Pivot tables in MS Access were used to compute species richness per grid cell. One query was conducted for each species group (Table 2.2).

2.2.2 ATKIS data

Information on the extent and spatial configuration of different land use types were derived from the ATKIS Digital Landscape Model (DLM, LGN, 2008) of Lower Saxony and Bremen. Overall, the DLM differentiates c. 90 different 'ATKIS object types'. These were combined into 8 different land cover types (LCTs), i. e. industrial, urban and traffic areas, croplands, forests, grasslands, water bodies and "other LCTs" as listed in Table 2.3. Linear elements represented as polylines in the DLM (roads, rivers) were buffered by using either standard values (roads, FGSV, 1996, see. Table A.5 in Appendix A) or information provided by the dataset itself (rivers) in order to convert them into polygons (Hawth's Analysis Tools, BEYER, 2004). Subsequently, they were combined with the other polygon features to a non-overlapping feature layer using ArcGIS-ArcInfo 9.2 (ESRI INC., 2007).

Information on average altitude, inclination and exposure per grid cell were derived from the ATKIS Digital Elevation Model (DEM, LGN, 2008). The DEM data were further used to calculate the *Mean Heat Load Index* (McCUNE & KEON, 2002, eq. 3) per grid cell by using the ArcScript 'Heat load index' (PARKS, 2004) in ArcView GIS 3.3 (ESRI INC., 2002).

Table 2.1: Overview of the data used in the analysis.

Description	Source	Data type & resolution	Derivable information
Species data			
Occurrence data of all 1879 vascular plant taxa of Lower Saxony and Bremen per grid cell	Vascular plant survey of Lower Saxony ^a , NLWKN ^b	MS Access data base (1993–2003), 2,395,045 entries	Species distribution, species richness
ATKIS data			
Digital Landscape Model (DLM)	LGN (2008), ATKIS	Shape file, scale 1:25,000	Landscape composition and configuration
Digital Elevation Model (DEM)	LGN (2008), ATKIS	Raster file, c. 5 m × 5 m	Altitude, exposure, inclination, heat load index
Areal information			
Geographical classification of natural landscapes (Naturräumliche Gliederung)	NLWKN, after PILGRIM & FRANKE, 1993	Shape file, scale 1:50,000	Natural landscape units, bio-geographical regions
Spatial information on protected areas (Natura 2000, nature protection areas, national parks, forest reserves)	NLWKN, 2010a; NLWKN, 2010b, NW-FVA ^c	Shape files, scale 1:25,000	Location and extent of protected areas
Environmental variables			
Map of soil types of Lower Saxony (BÜK 50 ^d)	LBEG ^e 2003	Shape file, scale 1:50,000	Soil types (27 categories), soil humidity (5 categories), CaCO ₃ content of the soil (3 categories), diversity of soil types
Climate data	DWD ^f 1961-1990	Raster files, 1 km × 1 km	Mean/min./max. annual temperature, mean/min./max. annual precipitation, no. of summer days, no. of days with frost/snow/ice, sunshine duration, phenological data

^a Niedersächsisches Pflanzenarteninventarprogramm, ^b Niedersächsisches Landesamt für Wasserwirtschaft, Küsten und Naturschutz, ^c Nordwestdeutsche Forstliche Versuchsanstalt, ^d Bodenübersichtskarte 1:50,000, ^e Landesamt für Bergbau, Energie und Geologie, ^f German Weather Service, ^g Deutscher Wetterdienst

Table 2.2: Groups of vascular plant species based on residence and threat status of the species as well as their habitat preferences as used in the analyses of the present thesis.

Levels	Definition
Threat status	
Threatened species	Species listed on the Red List of Lower Saxony and Bremen, categories 1, 2, 3, G, R (critically endangered, endangered, vulnerable, threatened, near threatened, extremely rare) ^a
Non-threatened species	Species not threatened according to the Red List of Lower Saxony and Bremen, categories *, V (not threatened, prewarned) ^a
Residence status	
Native species	Species native to Lower Saxony and Bremen ^b
Archaeophytes	Plant species not native to Lower Saxony and Bremen; established prior to 16 th century ^b
Neophytes	Plant species not native to Lower Saxony and Bremen; established in 16 th or later ^b
Habitat preference	
Forest species	Herbaceous and shrubby plant species linked to forest habitats ^c
Urban species	Plant species of urban habitats ^d
EIV species N1-3	Plant species of nutrient-poor habitats ^e
EIV species N7-9	Plant species of nutrient-rich habitats ^f

^a GARVE (2004), ^b Information taken from the data base of the vascular plant survey of Lower Saxony (NLDWKN 1993-2003), BfN (2013) and Klotz et al. (2002); ^c Categories K1.1, K1.2, S1.1, S1.2, Schmidt et al. (2011), ^d Urbanity indicator, categories 4 (urbanophilic) and 5 (moderately urbanophilic), Klotz et al. (2002), ^e Ellenberg Indicator 1, 2, 3 (BfN, 2013); ^f Ellenberg Indicator 7, 8, 9 (BfN, 2013).

2.2.3 Areal information

Information on bio-geographic regions and the location and extent of protected areas were obtained from shape files provided by the NLWKN and the NW-FVA (Table 2.1).

The geographical classification of natural landscapes of Lower Saxony (PILGRIM & FRANKE, 1993) defines the borders of the natural landscape units of the state. Roughly, the state area can be divided into three regions (coastal, lowland and upland area, cp. section 2.1) which themselves can be further divided into 10 bio-geographical regions and 82 natural landscape units (see Table A.1 in Appendix A). Because the grid cells in some cases were placed on the border between two or several regions/units the following rule was applied: Each grid cell was defined to belong to that region/unit that holds the highest proportion of area of the respective grid cell.

Information on protected areas included the location and extent of Natura 2000 sites, nature protected areas (*Naturschutzgebiete*, NSG), national parks as well as forest reserves (*Naturwaldreservate*) (Table 2.1).

2.2.4 Environmental variables

Soils Information on soil types, soil humidity, CaCO₃ content as well as the diversity of soil types were obtained from the map of soil types of Lower Saxony (BÜK 50, scale 1:50,000, vector data; Table 2.1). For all calculations the vector data were transformed into raster data of a pixel size of 50 m × 50 m.

BÜK 50 lists 126 soil types that were grouped into 27 soil type categories (see Table A.4 in Appendix A). Based on these categories, the diversity of soil types was calculated using Fragstats 3.3 (MCGARIGAL & CUSHMAN, 2002).

Soil humidity in BÜK 50 is arranged in 11 levels ranging from arid to wet/open water conditions. An additional 12th level includes all soils influenced by backwater. The 12 levels were grouped into 5 categories: "dry" (BÜK 50, soil moisture levels 0–3), "mesic" (soil moisture levels 4–6), "moist" (soil moisture levels 7–9), "wet" (soil moisture levels 10–11) and "influenced by backwater" (soil moisture level 12). Finally, the surface ratio of each soil moisture category was calculated per grid cell.

The CaCO₃ content of the soil in BÜK 50 is arranged in six levels ranging from soils with no CaCO₃ content (BÜK 50, level c0) to soils with a high CaCO₃ content (level c5). These levels were grouped into three categories: "free from CaCO₃" (BÜK 50 CaCO₃ content level c0, c1), "medium CaCO₃ content" (c2, c3) and "high CaCO₃ content" (c4, c5). Then, the surface ratio of each CaCO₃ category was calculated per grid cell.

Climate Information on regional climate were obtained from climate data of the German Weather service (*Deutscher Wetterdienst*, DWD; Table 2.1). Several different climatic parameters were derivable, i. e. mean, minimum and maximum annual temperature and precipitation, the number of summer days and days with frost, snow

Table 2.3: Land cover types (LCTs) derived from ATKIS-DLM. For more detailed information on the included ATKIS object types see Table A.6 in Appendix A.

LCT	Description
Industrial areas	Areas used for (non agricultural) production (e. g. supply and disposal facilities, exhibition centers, mining areas etc.)
Urban areas	Areas used for housing and recreation (e. g. residential areas, parks, campgrounds etc.)
Traffic areas	Areas used for transport (roads, railway lines, airports, etc.)
Croplands	Areas used for agricultural production
Forests	Forested areas
Grasslands	Grassland areas
Water bodies	Standing and flowing waters (e. g. lakes, streams, channels, springs)
Others	Areas not belonging to any of the above mentioned LCTs (e. g. tree rows, bogs, islands)

and ice as well as the sunshine duration. Furthermore, phenological data, e. g. the time of snowdrop (*Galanthus nivalis*) flowering, apple blossom etc. were available. All parameters were calculated as averages per grid cell.

2.2.5 Landscape metrics

Landscape metrics were used to quantify the composition and configuration of the landscape in the study area. These metrics are based on the patch-corridor-matrix concept by FORMAN (1995) which states that landscapes are composed of structural elements: the patches, the corridors and the matrix. The arrangement of these elements in the landscape creates a particular landscape pattern or patch mosaic (LANG & BLASCHKE, 2007). The term 'patch' in this context describes more or less homogeneous, non-linear landscape elements that differ from the surrounding landscape (LEITÃO et al., 2006). The matrix is defined by its proportion of the landscape ($\geq 50\%$), its connectivity as well as its control on landscape dynamics.

Landscape metrics that evolved in the 1980s and 90s (LANG & BLASCHKE, 2007) facilitate the description of patterns of a landscape with regard to its composition and configuration on the patch, class as well as landscape level. These indices describe and quantify the spatial structure of patches, classes and the landscape, respectively (LEITÃO et al., 2006). They provide the opportunity to measure geometric properties of landscape elements and their relative position and distribution in the landscape (LEITÃO et al., 2006). On the patch level, the geometric properties of individual patches can be assessed, whereas on the class level patches of the same type (class; in this study named land cover types, LCTs) are aggregated. On the landscape level the configuration of different types of classes is considered (LANG & BLASCHKE, 2007).

All landscape metrics were calculated using Fragstats 3.3 (MCGARIGAL & CUSHMAN, 2002). Because the software is limited to the use of raster data, the DLM vector

data were converted to raster format with a resolution of $5\text{ m} \times 5\text{ m}$ using ArcGIS-ArcInfo 9.2 (ESRI INC., 2007). A higher resolution could not be chosen because of hardware limitations. Because landscape metrics had to be assessed separately for each of the 1762 grid cells (see section 2.1) the raster data set was cut into 1762 separate rasters using Hawth's Tools (BEYER, 2004) in ArcGIS. In the following these 1762 tiles were considered as separate 'landscapes'. Landscape metrics were calculated on class and landscape level, respectively. The indices that were calculated describe the landscape composition (e. g. patch sizes, patch numbers) and configuration, i. e. the interactions and correlations between the landscape elements (e. g. expressed in terms of patch shape, complexity, fragmentation). The following paragraphs name and describe the landscape metrics used to quantify the composition and configuration of the landscape. Naming of the indices follows the nomenclature used in Fragstats 3.3 (McGARIGAL et al., 2002).

Landscape metrics quantifying landscape composition

Class level The landscape composition within the 1762 grid cells on the class level was assessed using the indices *Total Class Area* (CA) and *Number of Patches* (NP; Table 2.4). CA for each landscape reports the percentage of area of the particular classes (McGARIGAL & CUSHMAN, 2002). NP counts the number of patches of each class under observation. The NP index together with CA represents a simple measure to quantify fragmentation of the classes in the landscape: If a class is characterised by high values of CA as well as NP the area belonging to the class is highly fragmented.

Landscape level The landscape composition within the 1762 grid cells on the landscape level was assessed using the indices *Patch Richness* (PR), *Shannon's Diversity Index* (SHDI) and *Shannon's Evenness Index* (SHEI; Table 2.4). PR addresses the number of classes in the landscape and thus is an important basic measure for landscape diversity. If besides the pure number of classes in the landscape also their proportions shall be considered, the *Shannon's Diversity Index* is the method of choice: If the landscape is composed of one patch only, the value of SHDI equals zero. The more patch types are present and the more homogeneously distributed they are, the larger the value of SHDI becomes. SHEI, in turn, is a measure of the evenness of the distribution of patch types/classes (McGARIGAL & CUSHMAN, 2002). The index can take values between zero and one: If a landscape consists of only one patch type, values of SHEI will be zero. If several LCTs are present, the value of SHEI increases the more evenly distributed they are.

Table 2.4 lists all landscape metrics used to assess landscape composition, their range of values as well as their units.

Table 2.4: Landscape metrics used to assess landscape composition with respect to their range of values, units and basis of computation (after MCGARIGAL & CUSHMAN, 2002). LCT = land cover type; \uparrow = increase in value; n/a = not available.

Landscape parameter	Landscape metric	Range of values	Unit	Basis of computation
Class level				
Total class area	<i>Total Class Area, CA</i>	$CA > 0$	ha	All LCTs
Patch numbers/ Landscape diversity	<i>Number of Patches, NP</i>	$NP \geq 1$ $NP = 1$, if the landscape consists of one single patch	n/a	grassland, cropland, forest
Landscape level				
Number of patch types (classes)	<i>Patch Richness, PR</i>	$PR > 0$	n/a	All LCTs
Landscape diversity	<i>Shannon's Diversity Index, SHDI</i>	$SHDI \geq 0$ $SHDI = 0$, if the landscape consists of one patch/class only $SHDI \uparrow$, if the number of patch types/classes increases and/or the proportions of different classes is becoming more uniformly distributed	n/a	All LCTs
Uniform distribution of LCTs	<i>Shannon's Evenness Index, SHEI</i>	$0 \leq SHEI \leq 1$ $SHEI = 0$, if the landscape consists of one patch/class only $SHEI \uparrow$, if the proportion of the patches of all classes are becoming more equal $SHEI = 1$, if the patches of all classes are present at the same proportions	n/a	All LCTs

Landscape metrics quantifying landscape configuration

Class level Landscape configuration on the class level was assessed only for selected LCTs, i. e. forest, grassland and cropland areas. For these LCTs several configuration indices, namely the *Area-weighted Mean Patch Area Distribution* (AREA_AM), the *Area-weighted Mean Shape Index* (SHAPE_AM), the *Total Edge Index* (TE), the *Mean Proximity Index* (PROX_MN), the *Total Core Area Index* (TCA) as well as the *Area-weighted Mean Core Area Distribution* (CORE_AM) were calculated (Table 2.5). In the present study metrics based on area-weighted mean (AM) patch characteristics were favoured over those describing the average (MN) patch characteristics because the former provide a more landscape-centric instead of patch-centric perspective and thus better reflect the average conditions of a randomly chosen pixel in the landscape (McGARIGAL, 2014).

AREA_AM calculates the area-weighted mean size of all patches of one class. Thus, the index allows to discriminate between LCTs present at a large vs. small scale (McGARIGAL & CUSHMAN, 2002).

SHAPE_AM quantifies the complexity of patches. In raster data the simplest shape is a square; the more complex a patch becomes, i. e. the more it deviates from the shape of a square, the stronger the value of the *Shape Index* increases.

PROX_MN quantifies the isolation or rather connectivity of the patches of one class. For the calculation of the index a 'search radius' has to be defined; patches within this distance are regarded as connected. The search radius in the study was set to 100 m. If a patch has no neighbours of the same patch type within this distance the *Proximity Index* is 0; the index increases as the neighbourhood of the patch is increasingly occupied by patches of the same type; the closer these patches become and the less scattered they occur, the larger the value of the index becomes (McGARIGAL, 2014).

TE sums up the length of the edges of all patches belonging to one LCT (LANG & BLASCHKE, 2007). A high value of TE can – depending on the quality of the LCT under consideration – either be an indicator of a high structural diversity or identify a high degree of landscape fragmentation (LANG & BLASCHKE, 2007).

TCA sums up the core areas of all patches belonging to one class. The core area is defined as the area within a specified core area distance, i. e. that area that is not influenced by edge effects (McGARIGAL & CUSHMAN, 2002; LANG & BLASCHKE, 2007). The core area of patches, different from patch size, is influenced by the shape of the patches. Thus, a patch can be potentially large enough to support particular species but because of its shape can be influenced by edge effects to such a large extent that the habitat becomes unsuitable for species (McGARIGAL & CUSHMAN, 2002). For the calculation of TCA the 'edge depth' has to be defined. In the present study TCA was calculated for forests only; edge depth was defined as being 100 m (cp. MURCIA, 1995; HARPER et al., 2005; HENNENBERG et al., 2005; WEI & HOGANSON, 2005; HENNENBERG et al., 2008a,b). CORE_AM resembles TCA but measures the average core area of all patches belonging to one class.

Table 2.5: Landscape metrics used to assess landscape configuration with respect to their range of values, units and basis of computation (after MCGARIGAL & CUSHMAN, 2002). FGs = fragmentation geometries. LCT = land cover type; ↑ = increase in value; n/a = not available.

Landscape parameter	Landscape metric	Range of values	Unit	Basis of computation
Class level				
Patch complexity	<i>Shape Index Distribution, SHAPE_AM</i>	SHAPE_AM \geq 1 SHAPE_AM = 1, if the patches are maximally compact (square) SHAPE_AM ↑, if the shape of patches becomes more irregular	n/a	grassland, forest, cropland
Mean patch size	<i>Patch Area Distribution, AREA_AM</i>	AREA_AM > 0	ha	grassland, forest, cropland
Total Core Area	Total Core Area, TCA	TCA \geq 0 TCA = 0, if each location of all patches belonging to the class lays within the specified edge depth	ha	forest
Mean core area	<i>Core Area Distribution, CORE_AM</i>	CORE_AM > 0	ha	forest
Total edge length	<i>Total Edge, TE</i>	TE \geq 0	m	grassland, forest, cropland
Patch isolation	<i>Proximity Index, PROX_AM</i>	PROX \geq 0 PROX = 0, if a patch has no neighbours of the same patch type within the specified search radius	n/a	grassland, forest, cropland
Landscape level				
Complexity of patches	<i>Shape Index Distribution, SHAPE_AM</i>	SHAPE_AM \geq 1 SHAPE_AM = 1, if the patches are maximally compact (quadrat) SHAPE_AM ↑, if the shape of patches becomes more irregular	n/a	all LCTs
Mean patch size	<i>Patch Area Distribution, AREA_AM</i>	AREA_MN > 0	ha	all LCTs
Total edge length	<i>Total Edge, TE</i>	TE \geq 0	m	all LCTs
Landscape fragmentation	<i>Effective Mesh Size, MESH</i>	2.5×10^{-3} ha ^a \leq MESH \leq size of the grid cell	ha	FGs & remaining landscape

^a Area of one single pixel of the raster (pixel 5 m × 5 m = 25 m² = 2.5 × 10⁻³ ha).

Landscape level Landscape configuration on the landscape level was assessed using the *Area-weighted Mean Shape Index* (SHAPE_AM), the *Area-weighted Mean Patch Area Distribution* (AREA_AM) as well as the *Effective Mesh Size Index* (MESH; later following JAEGER et al. (2006) also named m_{eff}) (Table 2.5).

As explained for the class level, SHAPE_AM quantifies the complexity of patches. On the landscape level this complexity is not averaged over the patches of one class but over all patches being present in the landscape. The same applies to AREA_AM that on the landscape level calculates the area-weighted mean size of all patches in the landscape and, in doing so, provides a measure of the overall fragmentation of LCTs.

MESH measures the fragmentation of a landscape and is frequently used in many studies quantifying landscape fragmentation (e. g. JAEGER et al., 2001; WALZ, 2005; ES-SWEIN & SCHWARZ-VON RAUMER, 2006; MOSER et al., 2007; JAEGER et al., 2008; GIRVETZ et al., 2008; LI et al., 2010; EEA, 2011). The index denotes the size of the patches when the landscape is divided into S areas (each of the same size) with the same degree of landscape division as obtained for the observed cumulative area distribution (MCGARIGAL, 2014).

Table 2.5 lists all landscape metrics used to assess landscape configuration, their range of values as well as their units.

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3 | Patterns of vascular plant species richness at the landscape scale

Schmiedel, I.; Schacherer, A.; Hauck, M.; Schmidt, M. & Culmsee, H. (2011): Verbreitungsmuster der Farn- und Blütenpflanzen in Niedersachsen und Bremen unter Berücksichtigung ihres Einbürgerungsstatus und ihrer Gefährdungssituation. *Tuexenia* **31**: 211-226.

Summary

The present study analyses the richness patterns of different groups of vascular plant species for the federal states of Lower Saxony and Bremen. For this purpose, all 1819 established species occurring in the two states as well as different sub-groups of species were considered. Sub-groups were defined by using information on residence (1509 native species, 160 archaeophytes, 145 established neophytes) and threat status (643 threatened vs. 1176 non-threatened species) of the species, respectively. Because of their contrasting demands regarding their habitats it was assumed that species belonging to the various groups would occupy different ecological niches in the landscape; therefore – on a large scale – their centres of diversity were assumed to be spatially segregated from one another.

Data on species occurrences were obtained from the floristic mapping of the two states (plant survey of Lower Saxony, NLWKN 1982–2003) and were available on a grid basis (1,762 grid cells of c. 30 km²). The study aimed at answering the following questions: 1.) Which patterns of vascular plant species richness are observable in Lower Saxony and Bremen and the three natural landscape regions, respectively? 2.) Do the different plant species (sub-)groups show richness patterns that are distinguishable from each other and how much do their richness centers overlap? 3.) In which areas do threatened species show highest species numbers and how do these hotspots differ from that of non-threatened species?

Data were analysed by calculating the number of species per grid cell for each of the species groups mentioned above. Species richness per grid cell was then displayed using a geographical information system (GIS). To test if the centres of diversity of the species groups were overlapping, the pairwise non-parametric Spearman's rank correlation coefficient ρ was calculated.

Species richness of all groups was found to be heterogeneously distributed across the states with an observable increase of species numbers from north to south and – to some extent – also from west to east. The spatial distribution of the richness centres of the different plant species groups were largely correlated with each other but also showed some interesting divergences. Considering the total number of established species occurring in the study area an aggregated distribution pattern was observable: As site conditions in the Quaternary coastal and lowland areas are relatively homogenous these regions – with exception of the major stream valleys of the Weser, Aller and Elbe – were found to be relatively species poor. In contrast, the upland area with its heterogeneous site conditions, generally hosted a high overall species richness. The large group of native species showed a distribution very similar to that of the total number of species. Archaeophytes were concentrated in the coastal area of Bremen, the Weser and Aller plains, and the loess landscapes of the lowland area, as well as smaller cities. High richness of established neophytes was mainly found in congested urban areas. Richness centres of threatened species were spatially highly differentiated: at the coast only the islands in the North Sea repre-

sented small hotspots, while in the lowlands relatively large hotspots were situated in the Wendland, the Luneburg Heath and the Elbe-Weser triangle north of Bremen. In the uplands, the area around Göttingen, the uplands of the Weser and Leine valleys and the margins of the Harz Mountain represented well defined hotspots of Red List species.

Zusammenfassung

Voraussetzung für die Entwicklung von Schutzstrategien für den Pflanzenartenschutz ist die Kenntnis über die Verteilung der Zentren der Artenvielfalt im Raum. Je nach Einbürgerungsstatus und Gefährdungssituation kommt verschiedenen Artengruppen dabei eine unterschiedliche Bedeutung zu. In der vorliegenden Studie werden für die Gesamtfläche der Bundesländer Niedersachsen und Bremen die im Niedersächsischen Pflanzenarten-Erfassungsprogramm (1982-2003) auf Messtischblatt-Quadranten-Ebene erhobenen Verbreitungsdaten von Gefäßpflanzensippen unter Berücksichtigung der Gesamtlisfenliste (1.819 Sippen), ihres Einbürgerungsstatus (1.509 Indigene, 160 Archäophyten, 145 etablierte Neophyten) und ihrer Gefährdungssituation (ungefährdete und gefährdete Arten; davon 643 Sippen mit Rote-Liste-Status 1, 2, 3, G oder R) ausgewertet. Auf Basis der Gesamtlisfen ergibt sich eine inhomogene Verteilung der Sippendichte im Gesamtuntersuchungsraum, wobei die standörtlich relativ homogene Küste sowie das Tiefland – mit Ausnahme der großen Stromtäler (Weser, Aller, Elbe) – relativ artenarm sind und das standörtlich sehr heterogene Hugel- und Bergland grundsatzlich die hochsten Sippendichten aufweist. Unter Beruckichtigung des Einburgerungsstatus zeigen die Archaophyten jeweils die groten uberschneidungsbereiche zu den Indigenen und etablierten Neophyten. Die Verbreitungsmuster der groen Gruppe der Indigenen ahneln denen der Gesamtlisfen, wahrend sich die Archaophyten auf den Bremer Kustenraum, das Weser-Aller-Flachland, die Borden und das sudliche Weser-Leine-Bergland konzentrieren. Die Zentren der Sippenvielfalt der etablierten Neophyten liegen vor allem in stadtischen Ballungsraumen und erscheinen oftmals sehr punktuell. Die Rote-Liste-Arten sind in der Mehrzahl indigen (91%), 8% von ihnen sind Archa-, nur 1% Neophyten. Ihre Diversitatzentren sind auerordentlich differenziert: An der Kuste gehoren nur die isoliert liegenden Nordsee-Inseln dazu, wahrend im Tiefland das Wendland, die Luneburger Heide und das Elbe-Weser-Dreieck groflachige Diversitatzentren aufweisen. Im Hugel- und Bergland finden sich vor allem im Raum Gottingen, dem Weserbergland und am Harzrand gut abgegrenzte Zentren der Rote-Liste-Artendiversitat. Viele dieser bedrohten Sippen sind vermutlich Spezialisten, die an naturliche oder naturnahe Habitats angepasst und somit nur in den wenigen Landschaftsbereichen anzutreffen sind, die die entsprechenden Habitatbedingungen bieten.

Abstract

Distribution patterns of vascular plant taxa in the federal states of Lower Saxony and Bremen, Germany, with respect to their naturalisation and threat status

Detailed knowledge about the spatial distribution of richness centres is a prerequisite for developing national strategies for the conservation of vascular plant species. Thereby, different groups of plant taxa may have different relevance depending on their naturalisation and threat status. In this study, we investigate the spatial distribution of plant species richness centres in the area of the federal states of Lower Saxony and Bremen based on data from the plant survey of Lower Saxony (Niedersächsisches Pflanzenarten-Erfassungsprogramm). Analyses include: 1.) The complete floristic list (1819 taxa) 2.) three groups of taxa defined by their naturalisation status (1509 indigenous taxa, 160 archaeophytes, 145 established neophytes), 3.) two groups of taxa defined by their threat situation (unthreatened versus vulnerable taxa, including 643 taxa which have been red listed with status 1, 2, 3, G or R). Based on the complete floristic list, a clumped distribution pattern was observed. Site conditions in the Quaternary coastal and lowland areas are relatively homogenous and, with exception of the major stream valleys of the Weser, Aller and Elbe, relatively species poor, while the upland area, with its heterogeneous site conditions, shows the overall highest taxonomic richness. The large group of indigenous taxa shows a distribution very similar to that of the complete floristic list. Archaeophytes are concentrated in the coastal area of Bremen, the Weser and Aller plains, and the Old moraine loess landscapes, as well as smaller cities. High phytodiversity of established neophytes is mainly found in congested urban areas and several other small-scale centres of diversity. Red List species are mainly indigenous (91%), 8% are archaeophytes and only 1% are neophytes. Their centres of phytodiversity are spatially highly differentiated: at the coast only the islands in the North Sea are small hotspots, while in the lowlands relatively large hotspots are situated in the Wendland, the Luneburg Heath and the Elbe-Weser triangle north of Bremen. In the uplands, the area around Göttingen, the uplands of the Weser and Leine valleys and the Harz mountain margins are well defined hotspots of Red List taxa. Many of these endangered species are presumably specialists adapted and limited to natural and semi-natural habitats.

Keywords: Archaeophytes, plant diversity, phytodiversity, hotspots, neophytes, Red Lists, richness centres.

Einleitung

Das Verbreitungsmuster von Pflanzenarten im Raum wird durch deren Ausbreitungsmodi, ihr Konkurrenz- und Anpassungsverhalten, Wechselwirkungen mit der abiotischen Umwelt, durch evolutionäre und erdgeschichtliche, aber auch aktuelle, meist durch den Menschen verursachte Prozesse bestimmt (HAEUPLER, 1974). Landschaftsräume mit hoher struktureller und Habitat-Heterogenität, die sich aus einem Mosaik von künstlich geschaffenen Habitaten und Resten naturnaher Vegetation zusammensetzen, weisen oftmals eine sehr hohe Pflanzenartenvielfalt auf (DEUTSCHEWITZ et al., 2003). Insbesondere in städtischen Ballungszentren liegen die Artenzahlen meist höher als in dem umgebenden ländlichen Raum (HAEUPLER, 1974; WANIA et al., 2006; LIPPE & KOWARIK, 2008).

Für die Erfassung von Zentren der Artenvielfalt auf Landschaftsebene kommt verschiedenen Artengruppen je nach Einbürgerungsstatus und Gefährdungssituation eine unterschiedliche Bedeutung zu. Insbesondere Diversitätszentren von indigenen und/oder gefährdeten Pflanzensippen ("Hotspots") sind für den Artenschutz von Bedeutung (Globale Strategie zur Erhaltung der Pflanzen, SEKRETARIAT DER CBD, 2007), während solche von gebietsfremden Arten, insbesondere von invasiven Neophyten, auf eine Gefährdung von heimischen Arten, ökologischen Kreisläufen und des bestehenden Landschaftsbildes hinweisen können (HUBO et al., 2007). Gefährdete Arten stammen fast ausschließlich aus den Gruppen der Indigenen und Archäophyten (Definition bei WAGENITZ, 2003). In der mitteleuropäischen Landschaft sind die Hauptursachen für ihre Gefährdung die Intensivierung der Landnutzung und ihre Folgen (Eutrophierung, Entwässerung, Aufgabe extensiver Bewirtschaftungsweisen) sowie der gestiegene Flächenbedarf der Bevölkerung (GARVE, 2004). Etablierte Neophyten sind dagegen unter Einfluss des Menschen seit ca. 1500 n. Chr. in ein Gebiet eingewandert und – nach zeitlichen und populationsbiologischen Etablierungskriterien (SCHNITTLER & LUDWIG, 1996) – fester Bestandteil der Flora geworden. Aufgrund der kontrastierenden Habitatansprüche der verschiedenen Artengruppen ist anzunehmen, dass sie wie die heimischen Arten unterschiedliche Nischen in der Landschaft besetzen, so dass bei großräumiger Betrachtung ihre Zentren der Artenvielfalt räumlich getrennt sein sollten.

In dieser Studie werden für die Bundesländer Niedersachsen und Bremen die Verbreitungsmuster verschiedener Gefäßpflanzengruppen untersucht: Es werden die Gesamtheit der Arten sowie weitere, nach ihrem Einbürgerungsstatus (Indigene, Archäophyten und etablierte Neophyten) bzw. ihrer Gefährdungssituation (gefährdete und ungefährdete Arten) definierte Artengruppen betrachtet. Die Analyse umfasst die von 1982 bis 2003 erhobenen Funddaten von Sippen der Roten Liste und Florenliste der Farn- und Blütenpflanzen Niedersachsens und Bremens (GARVE, 2004) und basiert auf der floristischen Kartierung der beiden Bundesländer nach der Methodik des Niedersächsischen Pflanzenarten-Erfassungsprogramms (SCHACHERER, 2001). Dieses wird von der niedersächsischen Fachbehörde für Naturschutz (NLWKN) durchge-

führt und liefert auf Ebene von Messtischblatt-Quadranten flächendeckende Informationen über die Funde aller Gefäßpflanzensippen.

Die vorliegende Studie ist im Rahmen des von der Deutschen Bundesstiftung Umwelt geförderten Projektes "Identifizierung von Indikatorartengruppen für ein Biodiversitäts-Monitoring zur Bewertung von Grünland- und Waldlebensräumen" entstanden. Sie stellt eine grundlegende Auswertung des Pflanzenartendatensatzes dar, auf der weitere, speziellere Untersuchungen aufbauen. Auf Grundlage der Pflanzenartendaten sollen folgende Fragestellungen untersucht werden:

1. Welche Diversitätsmuster sind im gesamten Gebiet bzw. in den drei Naturräumen Küste, Tiefland sowie Hügel- und Bergland auf Grundlage der Gesamtliste der Farn- und Blütenpflanzen erkennbar?
2. Ergeben sich für die verschiedenen Gruppen je nach Einbürgerungsstatus unterschiedliche Muster der Phytodiversität und wie stark überschneiden sich die Zentren ihrer Diversität?
3. Wo befinden sich die Zentren der Sippenvielfalt der Rote-Liste-Arten (Hotspots im Sinne von SCHMITT & HAEUPLER, 2009) und wie unterscheiden sie sich von den Diversitätszentren der ungefährdeten Sippen?

Material und Methoden

Untersuchungsgebiet

Das Untersuchungsgebiet umfasst Niedersachsen und Bremen, wobei Niedersachsen mit einer Größe von 47.624 km² das zweitgrößte, Bremen mit einer Größe von 404 km² das kleinste Bundesland Deutschlands ist (NIEDERSACHSEN, 2008). Das Gebiet wird in drei Großregionen unterteilt: die Küste, das Tiefland und das Hügel- und Bergland (Abb. 3.1).

Die einzelnen Regionen unterscheiden sich stark in ihrer Naturraumausstattung: Die **Küste** wird geologisch vor allem durch marine Ablagerungen aus dem Holozän bestimmt (HEUNISCH et al., 2007). Ihr Klima ist ozeanisch geprägt, der mittlere Jahresniederschlag beträgt 700–900 mm, die mittlere Jahrestemperatur liegt bei durchschnittlich 9 °C. Vorherrschende Bodentypen sind Marschen, Syroseme, podsolige Ranker und Gleye (NLFB, 1997). Die Küstenregion wird dominiert von Grünländern, die etwa 57 % der Region einnehmen. Während Ackerflächen hier einen Anteil von 22 % besitzen, sind lediglich 1 % Wälder vorhanden.

Das niedersächsische **Tiefland** wird von pleistozänen Ablagerungen dominiert (HEUNISCH et al., 2007). Von West nach Ost bringt ein Kontinentalitätsgradient Jahresniederschlagsmittel von 800 mm im Westen bis etwa 550 mm im Osten sowie Jahresmitteltemperaturen von 8.0–8.5 °C mit sich. Entlang dieses Klimagradienten sind unterschiedliche Böden ausgebildet: Während im ozeanischen Westen von Sickerwasserbildung, Vernässung und Stoffverlagerung geprägte Böden (Moorböden, Podsole) vorherrschen, finden sich im subkontinentalen Osten vor allem Braunerden

und Ranker (NLFB, 1997). Das Tiefland wird dominiert von Ackerflächen (41 %), Wälder und Grünländer nehmen jeweils etwa knapp 1/4 der Region ein (21 bzw. 22 %).

Das standörtlich sehr heterogene **Hügel- und Bergland** wird vom Meso- und Paläozoikum (HEUNISCH et al., 2007) dominiert und weist aufgrund der stark variierenden Topographie besonders unterschiedliche klimatische Verhältnisse auf. Das Bergvorland ist geprägt von Löss- und Sandlössablagerungen sowie Ton-, Sand- und Kalkgesteinen (NLFB, 1997). Neben Parabraunerden und Braunerden kommen auch Übergänge zu Pseudogleyen vor (Bördenvorland). In der Lössbörde treten großflächig Schwarz- und Parabraunerden auf. Die Jahresdurchschnittstemperatur der tieferen Lagen (Becken) liegt 1–2 °C über der der Höhenzüge; die mittleren Jahresniederschläge sind auf den Höhenzügen deutlich höher (800–900 mm) als in den Beckenlagen (650–750 mm). Der Harz im Südosten Niedersachsens nimmt eine Sonderstellung ein: Der Hochharz (in Niedersachsen bis 971 m ü. NN) weist mittlere Jahresniederschläge von bis zu 1.500 mm und eine Jahresmitteltemperatur von nur 7 °C sowie lange, schneereiche Winter auf und ist somit (hoch-)montan geprägt (NLFB, 1997). Sowohl im Hochharz als auch im Oberharz, der submontane bis montane Klimaverhältnisse aufweist, kommen basenarme Silikatgesteine sowie Tonschiefer und Grauwacken vor. Vorherrschende Bodentypen sind hier Braunerden und Podsole (NLFB, 1997). Die Region weist von den drei Großregionen den höchsten prozentualen Waldanteil auf (32 %). Grünländer nehmen hier nur knapp 9 % der Fläche ein, während 45 % von Ackerflächen geprägt sind.

Untersuchungsraster

Das Untersuchungsgebiet ist auf Grundlage der Topographischen Karte 1:25.000 in ein Raster von Messtischblatt-Quadranten (MTB-Quadranten, im Folgenden nur Quadranten genannt) mit einer Größe von etwa 5.5 km × 5.5 km gegliedert. Für die vorliegenden Analysen wurden 1.762 Quadranten berücksichtigt, von diesen liegen 1.386 vollständig im Untersuchungsgebiet und sind gänzlich von Land bedeckt (Vollquadranten), weitere 376 liegen teilweise oder vollständig im Meer oder teilweise an der Grenze zu anderen Bundesländern (Randquadranten). Es wurden 255 Quadranten der Küste, 1.101 Quadranten dem Tiefland und 406 Quadranten dem Hügel- und Bergland zugeordnet. Dabei wurden Quadranten im Grenzgebiet zweier Regionen jeweils der Region zugeordnet, die den größeren Flächenanteil hatte.

Pflanzenartendaten

Die Gesamtliste der Farn- und Blütenpflanzen in Niedersachsen und Bremen richtet sich nach GARVE (2004). Verbreitungsangaben auf Quadranten-Ebene wurden der Datenbank des Niedersächsischen Pflanzenarten-Erfassungsprogramms entnommen (NLWKN 1982-2003, Stand 15.12.2008). Eine Sammlung von Artenverbreitungskar-

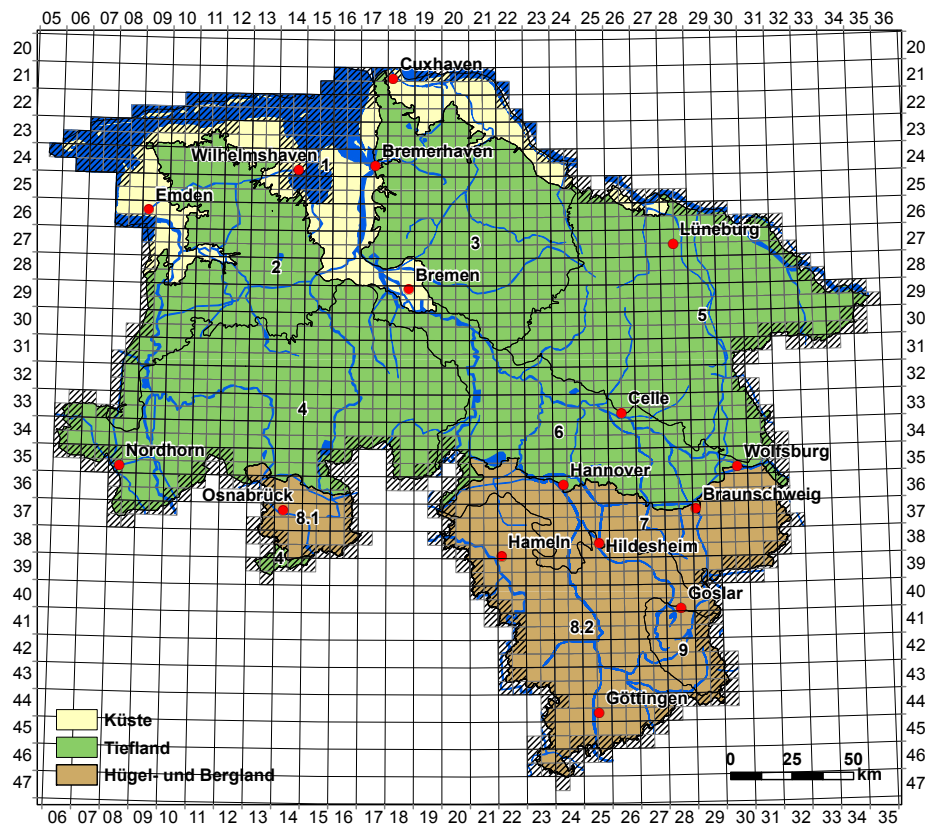


Abbildung 3.1: Das Untersuchungsgebiet umfasst die Gesamtfläche der Bundesländer Niedersachsen und Bremen und wird in drei naturräumliche Großregionen unterteilt: die Küste, das Tiefland und das Hügel-/Bergland. Eine weitere Untergliederung erfolgt in zehn naturräumliche Hauptregionen (1 = Watten und Marschen, 2 = Ostfriesisch-Oldenburgische Geest, 3 = Stader Geest, 4 = Ems-Hunte-Geest und Dümmer-Geestniederung, 5 = Lüneburger Heide und Wendland, 6 = Weser-Aller-Flachland, 7 = Börden, 8.1 = Osnabrücker Hügelland, 8.2 = Weser- und Leinebergland, 9 = Harz). Die 1.762 Messtischblatt-Quadranten sind grundlegende Kartiereinheit des Niedersächsischen Pflanzenarten-Erfassungsprogramms (Zahlenangaben am Rand entsprechen der TK25-Basisblatt-Nummer). Schraffierte Quadranten wurden auf einer kleineren Fläche floristisch erfasst, da sie teilweise oder gänzlich im Meer bzw. in angrenzenden Bundesländern liegen.

Figure 3.1: The study area the federal states of Lower Saxony and Bremen, with its three natural landscape regions (Küste, coast; Tiefland, lowlands; Hügel- und Bergland, uplands). The region can be further divided into ten natural regions (1 = Wadden Sea and marshes, 2 = East Frisian and Oldenburg moraine lowlands, 3 = Stade moraine lowlands, 4 = Ems, Hunte and Dümmer moraine lowlands, 5 = Lüneburg Heath and Wendland, 6 = Weser and Aller plains, 7 = Old moraine loess landscapes, 8.1 = Uplands of Osnabrück, 8.2 = Uplands of the Weser and Leine valleys, 9 = Harz mountains.) Plant occurrence data from the plant survey of Lower Saxony were available for 1762 quadrants of the topographical map (scale 1:25,000, i. e. about 5.5×5.5 km in size, frame numbers equal ordnance map numbers). The area of hatched quadrants has only been partly inventoried, because they are either only partially covered by land area or are situated in neighbouring federal states.

ten mit Angaben zur Funddichte wurde von GARVE (2007) veröffentlicht. Eine frühere Auswertung der Anzahl der Rote-Liste-Arten je Quadrant erfolgte durch GARVE (1994).

Der Pflanzenartendatensatz wurde nach folgenden Kriterien gruppiert:

1. Gesamtliste: alle Sippen der Farn- und Blütenpflanzen mit floristischem Normalstatus, d. h. Indigene, Archäophyten und etablierte Neophyten, unter Ausschluss von unbeständigen und synanthropen Vorkommen sowie Sippen, die in Niedersachsen und Bremen ausgestorben oder verschollen sind (1.819 Sippen mit Einträgen in der Datenbank).

2. Einbürgerungsstatus: Indigene (1.509 Sippen), Archäophyten (160 Sippen) bzw. etablierte Neophyten (145 Sippen) nach der von GARVE (2004) verwendeten Definition. Der Einbürgerungsstatus der Sippen wurde der Niedersächsischen Florenliste entnommen, wobei der Etablierungsstatus der Neophyten von GARVE (2004) übernommen wurde. Die Unterscheidung von Indigenen und Archäophyten erfolgte durch Abgleich mit der BIOLFLOR-Datenbank (KLOTZ et al., 2002). Für fünf der den Indigenen/Archäophyten zugeordneten Sippen konnte kein Einbürgerungsstatus festgestellt werden (*Panicum riparium*, *Persicaria mitis*, *Potamogeton x undulatus*, *Rumex x pratensis*, *Silene x hampeana*).

3. Gefährdungssituation: ungefährdete Sippen inklusive der Arten der Vorwarnliste (1.176 Sippen) bzw. gefährdete Sippen (643 Sippen mit Rote-Liste-Status 1, 2, 3, G oder R), wobei sich der Gefährdungsstatus auf das Gesamtgebiet Niedersachsen und Bremen bezieht (GARVE, 2004).

Datenanalyse

Für jede der untersuchten Gruppen wurde die Sippenzahl je Raumeinheit (Quadrant) erfasst und mithilfe eines Geographischen Informationssystems dargestellt (ArcGIS-ArcInfo 9.2; ESRI INC., 2007). Bei der Visualisierung der Verbreitungsmuster wurden natürliche Grenzwerte verwendet, wobei jeweils fünf Sippenzahl-Klassen gebildet wurden. In den Abbildungen wurden negative Abweichungen vom Mittelwert mit abgestuften Grautönen und die Klassen um bzw. oberhalb des Mittelwerts in Farbabstufungen dargestellt.

Zusätzlich wurden die mittleren und maximalen Sippenzahlen je Gruppe und Quadrant für Niedersachsen/Bremen bzw. die einzelnen Regionen Küste, Tiefland und Hügel- und Bergland ermittelt. Räumliche Autokorrelationen von Quadranten mit hohen Sippenzahlen (Klumpungen der Diversitätszentren) wurden als Moran's Index berechnet, wobei Werte gegen -1 auf eine starke Streuung und Werte gegen $+1$ auf eine starke Klumpung von Quadranten mit hoher Artenvielfalt im Raum hinweisen.

Für die nach ihrem floristischen Status (Indigene, Archäophyten, etablierte Neophyten) bzw. ihrem Gefährdungsstatus (ungefährdete Arten, Rote-Liste-Arten) gruppierten Sippen wurde jeweils für Paare dieser drei bzw. zwei Gruppen mittels des Rangkorrelationskoeffizienten ρ nach Spearman für nicht normalverteilte Daten

(*Shapiro-Wilk Normality-Tests*) mit der R-Software (R DEVELOPMENT CORE TEAM, 2012) überprüft, ob ihre Diversitätswerte (Zentren der Sippen-Diversität) miteinander korreliert sind. Der Koeffizient ρ kann Werte zwischen -1 und $+1$ annehmen, je nachdem, ob eine negative, positive bzw. – bei einer Annäherung gegen Null – keine Korrelation vorhanden ist.

Ergebnisse

Muster in der Phytodiversität der Gefäßpflanzen: Sippenzahlen auf Basis der gesamten Florenliste

Von den insgesamt 1.819 in Niedersachsen und Bremen vorkommenden Gefäßpflanzensippen mit floristischem Normalstatus kommen im Mittel etwa ein Viertel (435 Sippen) je Quadrant vor (Tab. 3.1). Maximal wurden 747 Sippen innerhalb eines Quadranten gezählt. Tabelle 3.2 listet die fünf Quadranten mit den höchsten Sippenzahlen auf Basis der gesamten Florenliste auf. Die sippenreichsten Quadranten kommen verstreut vor. Phytodiversitätszentren lassen sich im südlichen und östlichen Niedersachsen sowie nordwestlich von Bremen identifizieren (Abb. 3.2). Vor allem im nördlichen und westlichen Niedersachsen ist die Gefäßpflanzendichte gering. Die deutliche räumliche Konzentration von Diversitätszentren kann auch durch die starke räumliche Autokorrelation belegt werden (Tab. 3.3).

Vergleicht man die Gefäßpflanzenvielfalt der drei naturräumlichen Großregionen, so weist das Berg- und Hügelland sowohl die höchste absolute als auch die höchste mittlere Sippenzahl je Quadrant auf (Tab. 3.1). Die Sippenzahl ist in allen Quadranten relativ gleichmäßig hoch, eine Ausnahme stellt hier nur der Solling dar. Das Tiefland nimmt eine Mittelstellung ein: Die mittlere Sippenzahl je Quadrant entspricht in etwa dem für die Gesamtfläche ermittelten Durchschnitt. Während das westliche Tiefland relativ artenarm ist, finden sich Zentren der Sippen-Vielfalt vor allem im Weser-Aller-Flachland – insbesondere die Flusstäler von Aller, Leine und Weser zeichnen sich deutlich ab –, in der Lüneburger Heide und im Wendland. An der Küste kommen im Mittel nur 306 Sippen je Quadrant vor, so dass die Sippenzahl wesentlich unter dem Landesdurchschnitt liegt. Lediglich im Raum Bremen und auf den Inseln sind relativ hohe Sippenzahlen je Quadrant zu verzeichnen.

Table 3.1: Gesamtsippenzahl (Summe), mittlere Sippenzahl je Quadrant (MW) mit Standardabweichung (\pm Stabw) und maximale Sippenzahl pro Quadrant (Max) für verschiedene Artengruppen für Niedersachsen und Bremen (Gesamtfläche) bzw. die drei Regionen Küste, Tiefland und Hügel-/Bergland.

Table 3.1: Total number of vascular plant taxa (Summe), mean number of taxa per quadrant (MW) with standard deviation (\pm Stabw), and maximum number of taxa (Max) per quadrant for different plant groups considering the complete area of Lower Saxony and Bremen (Niedersachsen und Bremen), and the three regions coast (Küste), lowlands (Tiefland), and uplands (Hügel-/Bergland).

	Niedersachsen und Bremen			Küste			Tiefland			Hügel-/Bergland		
	Summe	MW \pm Stabw	Max	Summe	MW \pm Stabw	Max	Summe	MW \pm Stabw	Max	Summe	MW \pm Stabw	Max
Gesamtliste	1819	435 \pm 127	747	1133	306 \pm 152	651	1406	440 \pm 98	743	1484	503 \pm 119	747
Indigene	1509	347 \pm 101	611	888	239 \pm 116	482	1119	352 \pm 76	591	1206	401 \pm 97	611
Archäophyten	160	56 \pm 20	116	128	42 \pm 22	101	152	55 \pm 16	115	157	68 \pm 21	116
Etablierte Neophyten	145	32 \pm 13	84	115	25 \pm 16	80	132	33 \pm 11	79	119	34 \pm 13	84
Ungefährdete Arten	1176	408 \pm 114	668	935	292 \pm 142	601	1088	413 \pm 87	652	1017	469 \pm 104	668
Rote-Liste-Arten	643	27 \pm 19	127	266	14 \pm 13	62	437	27 \pm 17	111	526	34 \pm 23	127

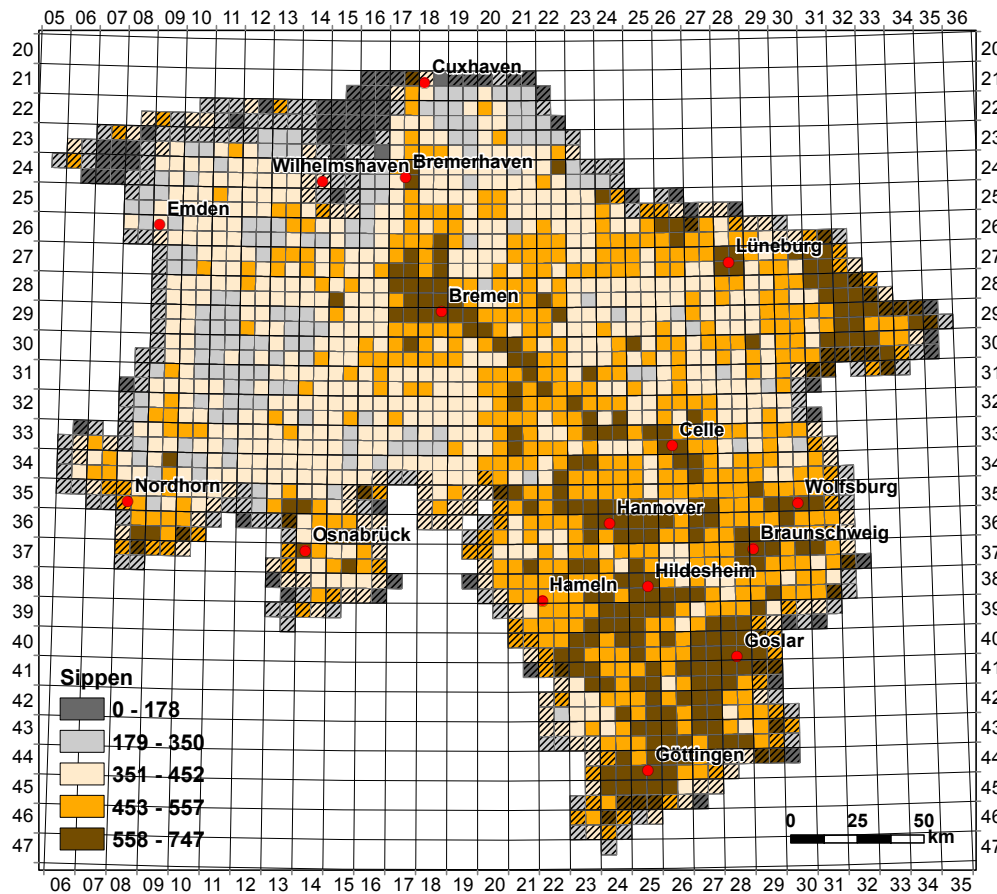


Abbildung 3.2: Anzahl der in Niedersachsen und Bremen vorkommenden Gefäßpflanzensippen mit floristischem Normalstatus je Quadrant basierend auf der Gesamtliste (1.819 Sippen).

Figure 3.2: Number of taxa (excluding erratic and synanthropic occurrences) per quadrant in Lower Saxony and Bremen, considering the complete list of 1819 vascular plant taxa.

Muster in der Phytodiversität der Gefäßpflanzen unter Berücksichtigung ihres Einbürgerungsstatus

Die absolute Zahl der Indigenen ist mit 1.509 Sippen um ein Zehnfaches höher als die der Archäophyten (160 Arten) und der etablierten Neophyten (145 Arten) (Tab. 3.1). Dementsprechend unterscheiden sich auch die mittleren Sippenzahlen zwischen den Gruppen mit durchschnittlich 347 Indigenen, 56 Archäophyten und 32 etablierten Neophyten je Quadrant. An der Küste kommen in allen Gruppen weniger Sippen vor als im Tiefland bzw. Hügel- und Bergland, wobei letztere Region – jeweils nur mit geringem Unterschied zum Tiefland – etwas artenreicher ist. Lediglich die absolute Zahl der etablierten Neophyten ist im Tiefland höher als in den übrigen Regionen.

Tabelle 3.2: Auflistung der jeweils fünf sippenreichsten Quadranten je Artengruppe . Quadrant = Nummer des Quadranten, Sippen = Anzahl der Sippen innerhalb des Quadranten, Ort = Innerhalb des Quadranten gelegene Stadt/ Ortschaft.

Table 3.2: A list of the five quadrants with the highest number of taxa per species group. Quadrant = Number of the quadrant, Sippen = Number of taxa inside the quadrant, Ort = City/Town located inside the quadrant.

Artengruppe		1	2	3	4	5
Gesamtliste	Quadrant	4028/3	2832/1	3729/2	4129/1	3032/3
	Sippen	747	743	736	727	726
	Ort	Langelsheim	Hitzacker	Weddel	Bad Harzburg	Luckau
Indigene	Quadrant	4429/2	4028/3	3032/3	4129/1	2832/1
	Sippen	611	598	591	590	586
	Ort	Walkenried	Langelsheim	Luckau	Bad Harzburg	Hitzacker
Archäophyten	Quadrant	3624/4	3624/1	3625/1	3729/1	3825/2
	Sippen	116	115	114	114	108
	Ort	Laatzen	Hannover	Misburg	Braunschweig	Hildesheim
Etablierte Neophyten	Quadrant	3729/1	2818/4	2918/2	3624/1	2818/3
	Sippen	84	80	80	79	76
	Ort	Braunschweig	Gröpelingen	Bremen	Hannover	Grambke
Ungefährdete Arten	Quadrant	3729/1	3729/2	4129/1	4028/3	3032/3
	Sippen	668	663	655	654	652
	Ort	Braunschweig	Weddel	Bad Harzburg	Langelsheim	Luckau
Rote-Liste-Arten	Quadrant	4429/2	2934/4	2832/1	4429/1	2731/4
	Sippen	127	111	110	109	108
	Ort	Walkenried	Gartow	Hitzacker	Bad Sachsa	Glienitz

Die Phytodiversitätskarte der indigenen Sippen (Abb. 3.3) weist starke Parallelen zur vorher besprochenen Darstellung der Sippen der gesamten Florenliste (Abb. 3.2) auf. Im Vergleich der drei Regionen ist die Sippenzahl je Quadrant im Hügel- und Bergland meist überdurchschnittlich hoch. Lediglich im Solling und in den Börden entsprechen die Sippenzahlen etwa dem Durchschnitt. Im Tiefland sind insbesondere weite Teile der Ems-Hunte-Geest und Dümmer-Geestniederung sehr artenarm. Das Wendland, Teile der Stader Geest und des Weser-Aller-Flachlandes – insbesondere wieder das Aller- und Leinetal – sind dagegen ausgesprochene Diversitätszentren. Mit Ausnahme des Raumes Bremen ist die Küstenregion nahezu flächendeckend arm an Indigenen. Die fünf sippenreichsten Quadranten (Tab. 3.2) sind anders gelagert als die der Gesamtliste, weisen jedoch wie diese ebenfalls eine recht zerstreute Verteilung auf.

Tabelle 3.3: Moran's Index als Maß für die räumliche Autokorrelation der Phytodiversitätszentren für die verschiedenen Gruppen.

Table 3.3: Moran's Index as a measure for spatial autocorrelation of plant richness centres considering all plant groups.

Artengruppe	Moran's Index	Z-Wert	p-Wert
Gesamtliste	0,138	148,68	0,01
Region Küste	0,157	26,04	0,01
Region Tiefland	0,110	75,25	0,01
Region Hügel-/Bergland	0,062	18,30	0,01
Indigene	0,133	143,18	0,01
Archäophyten	0,166	178,18	0,01
Etablierte Neophyten	0,090	97,69	0,01
Ungefährdete Arten	0,142	152,39	0,01
Rote-Liste-Arten	0,097	104,77	0,01

Ein differenzierteres Muster zeigt die Phytodiversitätskarte der Archäophyten (Abb. 3.4). Hier sind insbesondere der Bremer Küstenraum, das Weser-Aller-Flachland, die Börden sowie das südliche Weser-Leine-Bergland ausgesprochene Phytodiversitätszentren. Diese überschneiden sich stark mit denen der indigenen Sippen (Tab. 3.4), treten aber noch stärker räumlich konzentriert auf (Tab. 3.3). Dies zeigt sich auch bei der Betrachtung der fünf sippenreichsten Quadranten (Tab. 3.2): Die drei artenreichsten Quadranten konzentrieren sich im Raum Hannover, weiterhin ergeben sich hohe Sippenzahlen um Braunschweig und Hildesheim.

Die Phytodiversitätskarte der etablierten Neophyten (Abb. 3.5) zeigt zum einen eine großflächige Konzentration im Bremer Küstenraum, dem Weser-Aller-Flachland und den Börden, zum anderen aber auch eine Vielzahl von zum Teil sehr isolierten Einzelquadranten mit überdurchschnittlich hohen Sippenzahlen. Dabei sind städtische Ballungsräume (Großstädte ≥ 100.000 Einwohnern; HEINEBERG, 2006) grundsätzlich Zentren der Neophyten-Diversität. Diese starke Streuung von großflächigen und punktuellen Zentren der Neophyten-Diversität führt zu einem relativ geringen räumlichen Klumpungseffekt (Tab. 3.3). Dies spiegelt sich auch in der zerstreuten Verteilung der fünf sippenreichsten Quadranten wider (Tab. 3.2). Gleichzeitig ist jedoch der räumliche Überschneidungsbereich zwischen Archäophyten und etablierten Neophyten sehr groß und auch größer als der zwischen Indigenen und Archäophyten bzw. Indigenen und Neophyten (Tab. 3.4).

Muster in der Phytodiversität der Gefäßpflanzen unter Berücksichtigung ihrer Gefährdungssituation

Etwa ein Drittel der Gefäßpflanzen der Gesamtflorenliste (634 Sippen) sind als selten oder gefährdet eingestuft (Tab. 3.1). Für diese Gruppe werden im Hügel- und Berg-

Tabelle 3.4: Ähnlichkeiten in den Verbreitungsschwerpunkten der Gruppen nach Einbürgerungsstatus bzw. Gefährdungssituation. Ergebnisse der Berechnung des Rangkorrelationskoeffizienten ρ nach Spearman. ***, signifikant auf dem Niveau $p \leq 0,001$.

Table 3.4: Similarities in the distribution of species groups sorted by naturalization status (indigenous, archaeophytes, established neophytes) and threat situation (non-threatened vs. Red List taxa). Results based on the calculation of Spearman's Rank Correlation Coefficient ρ . ***, significant with $p \leq 0.001$.

	Spearman's ρ	p-Wert
Indigene - Archäophyten	0,744***	< 0,001
Indigene - etablierte Neophyten	0,681***	< 0,001
Archäophyten - etablierte Neophyten	0,805***	< 0,001
Rote-Liste-Arten - ungefährdete Sippen	0,691***	< 0,001

land sowohl die höchsten absoluten als auch mittleren Zahlen pro Quadrant erreicht. Die Mehrzahl der Rote-Liste-Arten gehört der Gruppe der Indigenen an, jedoch sind auch fast ein Drittel der Archäophyten Rote-Liste-Arten, während nur 6 % der Neophyten als gefährdet eingestuft werden (Tab. 3.5).

Die räumlichen Phytodiversitätsmuster der ungefährdeten Sippen (Abb. 3.6) weichen von denen der Rote-Liste-Arten (Abb. 3.7) stark ab, wobei der Überschneidungsbereich aber dennoch relativ hoch bleibt (Tab. 3.4). Insgesamt sind die Rote-Liste-Arten stärker auf mehrere, kleinflächigere Raumeinheiten verstreut als die stark zusammenhängenden Zentren der ungefährdeten Arten (Tab. 3.3). In der insgesamt eher artenarmen Küstenregion bilden nur die Nordsee-Inseln isolierte Hotspots der Rote-Liste-Arten (zum Hotspot-Begriff siehe SCHMITT & HAEUPLER, 2009). Im Tiefland ist die Überlagerung von Verbreitungsmustern der ungefährdeten und gefährdeten Sippen relativ hoch, wobei insbesondere das Wendland (vgl. Tab. 3.2), die Lüneburger Heide und das Elbe-Weser-Dreieck nördlich von Bremen Hotspots der Rote-Liste-Arten sind. Während im Hügel- und Bergland die Anzahl der ungefährdeten Sippen fast überall überdurchschnittlich hoch ist, weisen die Rote-Liste-Arten dagegen stark abgegrenzte Hotspots im Raum Göttingen, dem Weserbergland und am Harzrand (vgl. Tab. 3.2) auf.

Diskussion

Die räumliche Verteilung der Pflanzensippenvielfalt in Niedersachsen und Bremen weist für alle betrachteten Gruppen sowohl von Nord nach Süd als auch in gewissem Maße von West nach Ost eine Zunahme in der Sippendichte auf. Dabei ist das Hügel- und Bergland grundsätzlich die Region mit der höchsten Sippendichte. Eine Ausnahme bilden hier nur die Neophyten.

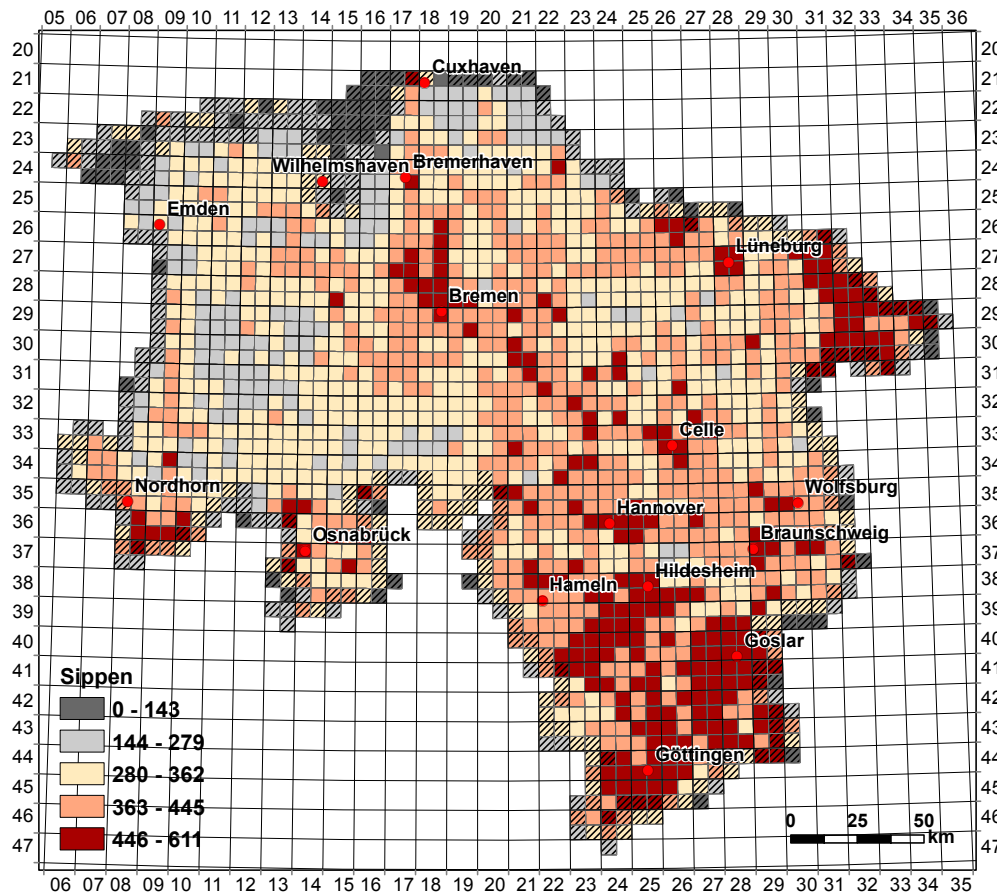


Abbildung 3.3: Anzahl der in Niedersachsen und Bremen vorkommenden 1.509 indigenen Gefäßpflanzensippen.

Figure 3.3: Number of taxa per quadrant in Lower Saxony and Bremen, considering 1509 indigenous vascular plant taxa.

Die relative Artenarmut des norddeutschen Tieflandes ist sicherlich u. a. dadurch bedingt, dass die landschaftliche Heterogenität (Klima, Relief, Gesteine, Böden) des Hügel- und Berglandes von den Lössböden bis in den Hochharz wesentlich höher ist als die des relativ uniform vom Quartär geprägten Tieflandes. Die kleinräumigen ökologischen Gradienten dürften somit zusätzlich sowohl eine höhere absolute Sippenzahl als auch höhere Sippendichten im südlichen Niedersachsen bedingen. Das insbesondere im Tiefland ausgeprägte Ost-West-Gefälle verläuft parallel zum klimatischen subkontinental-ozeanischen Gradienten. Insbesondere im nördlichen Harzvorland und im Wendland liegen die westlichen Verbreitungsgrenzen vieler Gefäßpflanzenarten, die Vorposten einer östlich der Elbe reicheren, subkontinental getönten Flora sind.

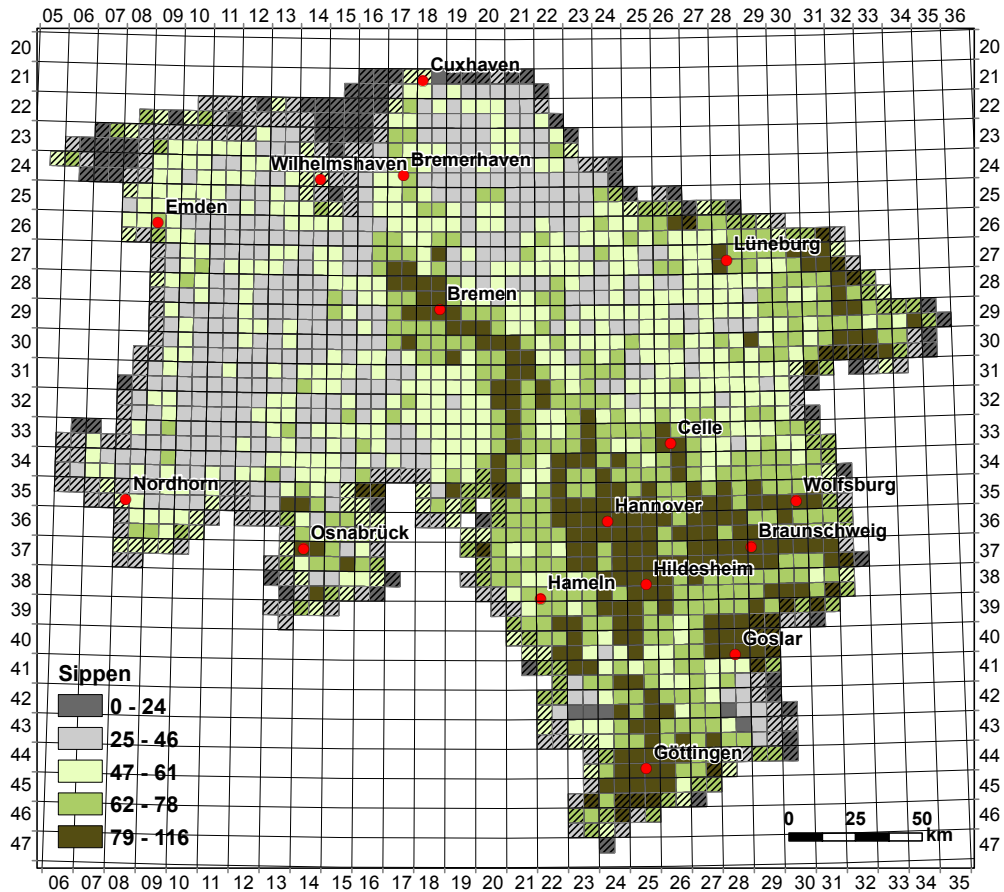


Abbildung 3.4: Sippenzahl je Quadrant der in Niedersachsen und Bremen vorkommenden 160 Archäophyten.

Figure 3.4: Number of taxa per quadrant in Lower Saxony and Bremen, considering 160 archaeophytes.

Die Verbreitungsmuster der Artengruppen nach Einbürgerungsstatus weisen große räumliche Überschneidungen auf. Während die Lage der Diversitätszentren von Indigenen und etablierten Neophyten im Tiefland viele Parallelen aufweist, sind sie im Hügel- und Bergland überwiegend negativ korreliert. KÜHN et al. (2003) zeigten in einer deutschlandweiten Analyse, dass die geologische Heterogenität eines Gebietes der wichtigste Faktor für die Sippenzahlen sowohl der Einheimischen als auch der Gebietsfremden ist. PYŠEK et al. (2002) erklärten die positive Korrelation von Indigenen und Neophyten in Naturschutzgebieten Tschechiens mit einer hohen Habitatdiversität in den entsprechenden Gebieten, die dazu führt, dass die beiden Gruppen nicht unbedingt miteinander in Konkurrenz stehen müssen. Dies mag auch für die positiv korrelierenden Vorkommen im niedersächsischen Tiefland zutreffen. Gerade in den Gebieten des Hügel- und Berglandes, in denen große, zusammenhängen-

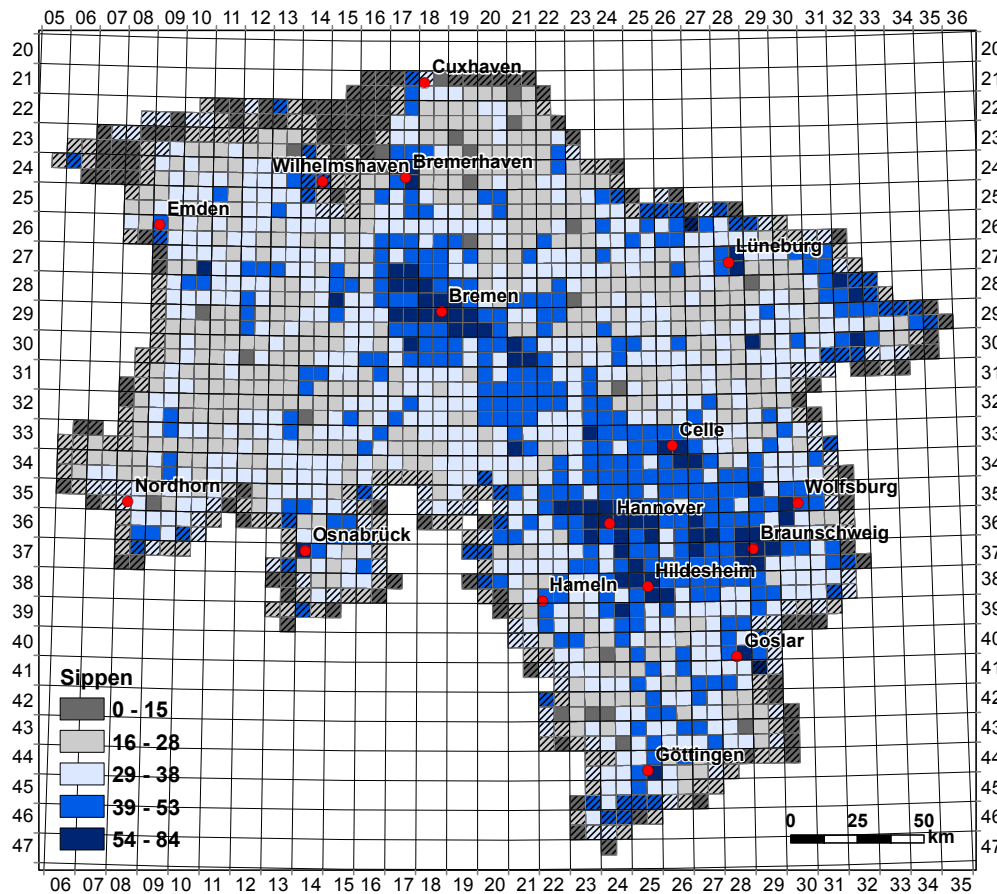


Abbildung 3.5: Sippenzahl je Quadrant der in Niedersachsen und Bremen vorkommenden 145 etablierten Neophyten.

Figure 3.5: Number of taxa per quadrant in Lower Saxony and Bremen, considering 145 established neophytes.

de Zentren der Sippendiversität der Indigenen und besonders geringe Neophyten-Dichten auftreten (Weserbergland, Göttinger Wald, Harz), sind jedoch auch noch besonders große, relativ zusammenhängende Waldflächen vorhanden. Natürliche und naturnahe Wälder, die in Mitteleuropa flächendeckend die potentiell natürliche Vegetationsformation darstellen (BOHN et al., 2000/2003), sind ein Refugium für eine Vielzahl der heimischen Arten. Dagegen ist die Anzahl der Neophyten in Wäldern geringer als im Offenland. Weniger als ein Drittel der in Niedersachsen etablierten Neophyten kommen nach SCHMIDT et al. (2003) regelmäßig in Wäldern vor. Ihre Schwerpunkte in bewaldeten Landschaften liegen auf Kahlschlagflächen und in bach- und wegbegleitenden Vegetationstypen (vgl. CHYTRÝ et al., 2005).

Besonders hoch ist die Übereinstimmung der Verbreitung der Archäophyten mit der von etablierten Neophyten. Gemeinsam ist beiden Artengruppen, dass sie im

Tabelle 3.5: Zahl der Gefäßpflanzensippen nach ihrem Einbürgerungsstatus (Indigene, Archäophyten und Neophyten) und ihrer jeweiligen Gefährdungssituation (ungefährdete und Rote-Liste-Arten) auf Grundlage der Gesamtflorenliste für Niedersachsen und Bremen.

Table 3.5: Vascular plant taxa richness sorted by naturalization status (indigenous, archaeophytes, established neophytes) and threat situation (non-threatened vs. Red List taxa) based on the complete floristic list of Lower Saxony and Bremen.

	Gesamtliste	Ungefährdete Sippen	Rote-Liste-Arten
Indigene	1509	925	584
Archäophyten	160	111	49
Etablierte Neophyten	145	136	9
Sippen ohne floristischen Status	5	4	1

Gefolge des Menschen nach Mitteleuropa eingewandert sind. Die meisten Archäophyten sind in Acker-, Grünland- und Ruderalvegetation angesiedelt. Sie sind damit zum Teil wichtige Bestandteile von historisch bedeutsamen, heute gefährdeten Biototypen, die relativ kleinflächig in eine ansonsten intensiv genutzte Kulturlandschaft eingebettet sind. Dies zeigt sich auch an der Vielzahl der als gefährdet eingestuft Archäophyten. Der von KÜHN et al. (2003) festgestellte hohe Anteil von Archäophyten in Lössgebieten stimmt in der vorliegenden Auswertung mit einem Schwerpunkt der Vorkommen dieser Artengruppe in den Lössböden überein. Letztere sind seit langem besiedelte und heute intensiv genutzte Agrar- und Industrielandschaften, in denen sich auch Neophyten großflächig etablieren konnten.

Urbane Ballungsräume weisen in Niedersachsen und Bremen generell eine hohe Artenzahl auf. Dies stimmt mit den Ergebnissen anderer Untersuchungen, etwa denen von HAEUPLER (1974), WANIA et al. (2006) und LIPPE & KOWARIK (2008) überein, die feststellten, dass urbane Lebensräume generell eine höhere Pflanzenartenzahl aufweisen als ländliche Gebiete gleicher Größe. Sie führten dies auf die hohe Landschaftsheterogenität in den vielfältig genutzten Räumen zurück, die in der Regel ein kleinräumiges Mosaik von standörtlich sehr heterogenen naturnahen und künstlich geschaffenen Lebensräumen darstellen und somit einer Vielzahl von Pflanzenarten als Wuchsorte dienen können. KÜHN et al. (2003) stellten außerdem fest, dass die Flora deutscher Städte natürlicherweise artenreich ist. Sie begründeten dies damit, dass Städte oftmals in von vorneherein sehr artenreichen Landschaftsbereichen entstanden sind.

Besonders evident ist der Zusammenhang zwischen der Diversität und der Urbanität der untersuchten Raumeinheiten für die Neophyten. Zu ähnlichen Ergebnissen kamen auch andere Studien, die sich mit der Verbreitung dieser Artengruppe beschäftigen (PYŠEK, 1998; ROY et al., 1999; HONNAY et al., 2003; CHYTRÝ et al., 2005; MASKELL et al., 2006; BOTHAM et al., 2009; NOBIS et al., 2009). MASKELL et al. (2006) wiesen für den ländlichen Raum Großbritanniens nach, dass nichtheimische Arten sich vor allem

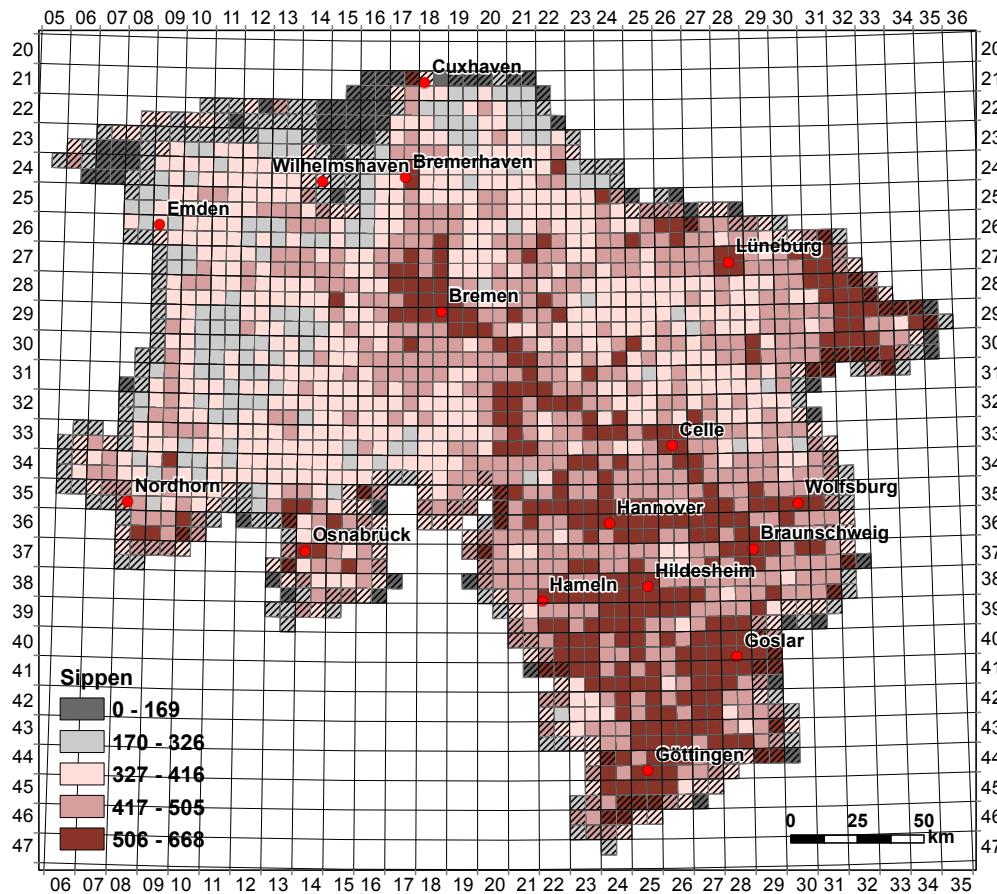


Abbildung 3.6: Anzahl der in Niedersachsen und Bremen je Quadrant vorkommenden 1176 ungefährdeten Gefäßpflanzensippen.

Figure 3.6: Number of taxa per quadrant in Lower Saxony and Bremen, considering 1176 invulnerable vascular plant taxa.

in vom Menschen geprägten Habitaten etablierten. BOTHAM et al. (2009) untersuchten in Großbritannien innerhalb von zwei verschiedenen Untersuchungszeiträumen (1987-1988 und 2003-2004) die Bindung von indigenen Pflanzenarten, Archäophyten und Neophyten an verschiedene Landbedeckungsklassen. Sie stellten fest, dass Neophyten in beiden Zeitabschnitten stark an urbane Landschaftsbereiche gebunden waren, wohingegen die Archäophyten nur im ersten Untersuchungszeitraum stark mit urbanen Lebensräumen assoziiert waren. In geringerem Maße als für die Neophyten ist für die Archäophyten auch in unserer Studie ein Zusammenhang zwischen Urbanität und Artenreichtum zu erkennen. Diese Gruppe zeigt relativ kleinflächig ausgeprägte Diversitätszentren im Umfeld vieler niedersächsischer Städte. Aufgrund der Großräumigkeit der zugrundeliegenden Raumeinheiten konnte hier nicht untersucht werden, ob der Artenreichtum der Archäophyten, wie PÝŠEK (1998) bemerkt,

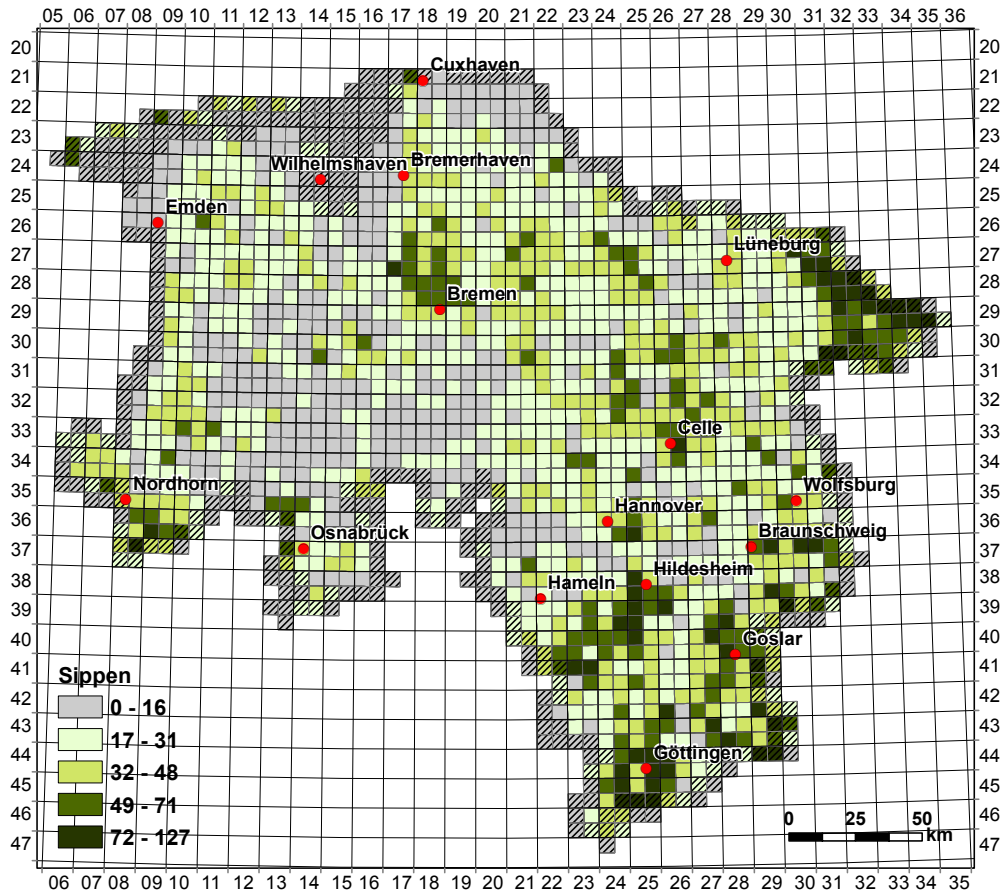


Abbildung 3.7: Anzahl der 643 seltenen und gefährdeten Gefäßpflanzensippen Niedersachsens und Bremens (Rote Liste-Status 1, 2, 3, G, R) je Quadrant.

Figure 3.7: Number of taxa per quadrant in Lower Saxony and Bremen, considering 643 rare and endangered vascular plant taxa (Red List status 1, 2, 3, G, R).

aufgrund ihrer Anpassung an ländliche Räume vor allem auch in kleineren Städten erhöht ist.

HAEUPLER & VOGEL (1999) sowie HAEUPLER (2000) betonen, dass die Musteranalyse von Rote-Liste-Artanzahlen gut geeignet ist, die 'echten' Hotspots der Gefäßpflanzen-diversität aufzudecken. Für Niedersachsen und Bremen weisen die gefährdeten Arten ein wesentlich differenzierteres und eher punktuelleres Verbreitungsmuster auf als die anderen betrachteten Artengruppen. Die Hotspots der Rote-Liste-Arten korrelieren in vielen Fällen mit denen der anderen Artengruppen, liegen jedoch nicht unmittelbar in den städtischen Ballungszentren. Bei den Rote-Liste-Arten handelt es sich fast ausschließlich um Indigene bzw. Archäophyten, wobei diese Arten oft Spezialisten sind, die an natürliche oder naturnahe Habitate angepasst sind. Eine besonders starke Überschneidung von Raumeinheiten mit hoher Dichte sowohl gefährdeter Arten

als auch von Neophyten ergeben sich in Niedersachsen für das Wendland, die Umgebung von Göttingen, Hildesheim und Goslar sowie die Region nördlich von Bremen. Es kann jedoch davon ausgegangen werden, dass die Hauptursachen der Gefährdung von Pflanzenarten in der Intensivierung der Landnutzung und ihren Folgen (Eutrophierung, Entwässerung, Aufgabe von Extensivflächen) sowie dem gesteigerten Flächenbedarf der Bevölkerung liegen und nicht durch die Konkurrenz der Neophyten bedingt sind (GARVE, 2004).

Danksagung

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4

The effectiveness of protected areas for the conservation of threatened vascular plant species

Schmiedel, I.; Schmidt, M.; Schacherer, A. & Culmsee, H. (2013): Die Effektivität von Schutzgebieten für die Erhaltung seltener und gefährdeter Gefäßpflanzenarten. Eine Untersuchung im niedersächsischen Tiefland. *Naturschutz und Landschaftsplanung* 45: 45-52.

Summary

The present study tests the effectiveness of protected areas for the conservation of rare and threatened vascular plant species. For this purpose, the lowland area of the federal state of Lower Saxony, Germany with its 41 geographical sub-regions served as a model region.

The analysis was conducted for nature protected areas¹ and Natura 2000 sites that represent the most important and extensive conservation areas in the lowlands. Information on the occurrences of Red List (RL) species were obtained from the floristic mapping of the state (plant survey of Lower Saxony, NLWKN 1982-2003). In addition to the total number of RL species, those species linked to forest habitats² (forest affinity categories 1.1 and 1.2, SCHMIDT et al., 2011) were considered in the analysis. All data were available on the basis of a regular grid of 15.729 grid cells with a size of 207 ± 2 ha. The study aimed at answering the following research questions: 1.) What patterns of diversity can be observed for the two species groups? 2.) Are occurrences of threatened species covered by nature protected areas? 3.) Does the designation of Natura 2000 sites improve the situation? 4.) Are the results obtained for RL species transferable to RL forest species?

The correlation between the extent of protected areas and the number of RL species per grid cell was analysed using Generalised Linear Models (GLM) with Poisson distribution. The calculation was performed separately for each of the 41 geographical sub-regions. The same kind of analysis was repeated for RL forest species.

Richness of the rare and threatened vascular plant species in the lowlands of Lower Saxony showed a distinct pattern. Centres of species richness (hotspots) were identified that should be considered as priority areas for nature conservation. Hotspots of RL species were mainly located in the Wendland area, the Westmünsterland and the Wesermünder Geest, those of RL forest species in the Oldenburger Geest. In contrast, large parts of the western lowlands as e. g. the Hunte-Leda-Moorniederung, the Bersenbrücker Land as well as the Bourtangener Moor were found to be very species-poor. With the Ostmünsterland and the Rahden-Diepenauer Geest being an exception, hotspots of RL species were well covered by protected areas. In contrast, richness centres of RL forest species were effectively protected in only c. 60 % of the geographical sub-regions. However, the number of species in those regions where no positive correlation between the distribution of species richness and the extent of protected areas could be found was generally very low. By including Natura 2000 sites the proportion of well-protected hotspots increased to an effective protection in 95 % of the geographical sub-regions for RL species and 81 % for RL forest species.

The diverging results obtained for the two species groups suggest that it is reasonable to not only focus on the total number of RL species representing a wide

¹German: *Naturschutzgebiete*

²Below named 'RL forest species'.

ecological amplitude as it has been done in most studies dealing with the effectiveness of protected areas for species conservation. Instead it is worthwhile to extend the analysis to species groups that represent more narrow habitat requirements.

Abstracts

Die Einrichtung von Schutzgebieten ist ein zentrales Instrument des Artenschutzes. Am Beispiel der 41 naturräumlichen Regionen des niedersächsischen Tieflands wird ein Verfahren zur Überprüfung der Effektivität von Naturschutz- (NSG) und FFH-Gebieten (Fauna Flora Habitat) für den Schutz seltener und gefährdeter Gefäßpflanzenarten (Rote-Liste-Arten = RL-Arten) vorgestellt. Grundlage sind die Daten des Niedersächsischen Pflanzenarten-Erfassungsprogramms.

In 85 % der naturräumlichen Regionen ist die Gesamtzahl der RL-Arten durch NSG effektiv geschützt; für die an den Lebensraum Wald gebundenen RL-Arten hingegen trifft das nur in 61 % der naturräumlichen Regionen zu. Die Ausweitung des Schutzgebietssystems durch FFH-Gebiete verbessert die Schutzsituation der RL-Arten insgesamt wie auch die der RL-Waldarten erheblich.

Der vorgestellte Ansatz ermöglicht die Identifizierung von Zentren der Artenvielfalt (Hotspots), die in Hinsicht auf ihren Schutz und ggf. gezielte konservierende Pflegemaßnahmen besondere Beachtung finden sollten.

Effectiveness of protection areas for the conservation of rare and endangered vascular plant species – An investigation in the lowlands of Lower Saxony, Germany

The establishment of protected areas is a central tool in species conservation. Using the 41 geographical regions of the Lowlands of Lower Saxony, Germany, as a model region, a method to test the effectiveness of nature protected areas and Natura 2000 sites for the conservation of rare and endangered vascular plant species (RL, Red List species) is proposed. RL species are effectively protected by nature protected areas in 85 % of the geographical regions; for RL species linked to forest habitats this is the case only in 61 % of the geographical regions. The extension of the protected area system to include Natura 2000 sites greatly improves the situation.

The approach allows the identification of regions with high species richness of endangered plants (hotspots) that should be particularly considered for conservation and management actions.

Einleitung

Die Einrichtung von Schutzgebieten stellt ein zentrales Instrument von Strategien zur Verringerung von Biodiversitätsverlusten dar (MEA, 2005; GASTON et al., 2006; BMU, 2007). In diesem Zusammenhang fordert die 'Globale Strategie zur Erhaltung der Pflanzen' (GSPC, CBD-COP6, Entscheidung VI/9, SEKRETARIAT DER CBD, 2007) einen effektiven in-situ-Schutz, durch den dem weiteren Rückgang von gefährdeten Pflanzenarten entgegengewirkt werden soll. Ob die für den Pflanzenartenschutz bedeutsamen Flächen auch tatsächlich durch Schutzgebiete abgedeckt sind, wurde bisher für größere Räume kaum untersucht. Für Deutschland fehlen Studien, die die Wirksamkeit der bestehenden Schutzgebietskulisse analysieren (ENGEL et al., 2012), und auch auf internationaler Ebene sind nur wenige Untersuchungen zur Effektivität von Schutzgebieten für den Artenschutz bekannt (DIMITRAKOPOULOS et al., 2004; DEGUISE & KERR, 2006; JACKSON et al., 2009).

Naturschutzgebiete (NSG) und Fauna-Flora-Habitat-Gebiete (FFH-Gebiete) des europäischen Netzwerks Natura 2000 sind in Deutschland die flächenmäßig bedeutsamsten Schutzgebietskategorien (BfN, 2008). Die Ausweisung von NSG kann aus wissenschaftlichen, naturgeschichtlichen und/oder landeskundlichen Gründen sowie wegen der Seltenheit, besonderen Eigenart oder hervorragenden Schönheit der Gebiete erfolgen (§ 23 BNatSchG, 2009). Der Artenschutz ist also nicht unbedingt explizites Kriterium für die Schutzgebietsausweisung. Im Rahmen der FFH-Richtlinie (Richtlinie 92/43/EWG) werden Schutzgebiete dagegen vorrangig nach Kriterien des Habitat- und Artenschutzes ausgewiesen (SSYMANK, 1998). Hierbei wurden allerdings vielfach bestehende Schutzgebiete, insbesondere NSG, eingeschlossen. Für beide Schutzgebietskategorien gibt es keine allgemeingültigen Schutzbestimmungen – diese müssen durch Schutzgebietsverordnungen (NSG) bzw. Managementpläne (FFH-Gebiete) festgelegt werden.

Die vorliegende Studie untersucht die Effektivität von Schutzgebietssystemen (NSG und FFH-Gebiete) für die Erhaltung von seltenen und gefährdeten Gefäßpflanzen im niedersächsischen Tiefland. Dabei werden zunächst alle im niedersächsischen Tiefland vorkommenden Gefäßpflanzenarten der Roten Liste Niedersachsens betrachtet (490 Sippen; Rote Liste-Status der Tiefland-Region 1, 2, 3, G, R; GARVE, 2004; im Folgenden RL-Arten genannt).

Um feststellen zu können, ob die für die RL-Arten der Gesamtliste erhaltenen Ergebnisse gleichermaßen für einen Teil der an einen bestimmten Lebensraum gebundenen RL-Arten gelten, werden die Analysen beispielhaft für die seltenen und gefährdeten Waldarten (nach SCHMIDT et al., 2011; 60 Sippen; im Folgenden RL-Waldarten genannt) wiederholt, da die Datenlage zu dieser Artengruppe sehr gut ist und Waldlebensräume eine große ökologische Bedeutung besitzen (ML, 2004). Wälder, insbesondere solche mit langer Habitatkontinuität, nehmen im nordwestdeutschen Tiefland im Vergleich zum nordostdeutschen Tiefland und vielen deutschen Mittelgebirgsregionen einen sehr geringen Flächenanteil ein (GLASER & HAUKE, 2004; SCHMIDT

et al., 2009). Hinsichtlich der Nutzungsgeschichte und Größe der Wälder sowie ihres Fragmentierungsgrades bestehen zwischen den 41 naturräumlichen Regionen des Tieflands große Unterschiede, die sich sowohl in der Ausstattung mit seltenen und gefährdeten Waldgefäßpflanzenarten als auch in der Schutzgebietskulisse widerspiegeln sollten.

Die NSG und FFH-Gebiete im niedersächsischen Tiefland sind sehr unterschiedlichen Alters. Während die NSG teilweise bereits seit über 90 Jahren bestehen (POHL, 1999), wurden die FFH-Gebiete als Teile des Natura-2000-Schutzgebietssystems erst seit 1997 an die EU gemeldet (NMUK, 2010).

Folgenden Fragestellungen wird nachgegangen:

1. Wie ist die Artenvielfalt der betrachteten Artengruppen im niedersächsischen Tiefland verteilt?
2. Inwieweit sind die Vorkommen der RL-Arten durch NSG abgedeckt?
3. Hat sich die Schutzgebietsabdeckung durch die Ausweisung von FFH-Gebieten verbessert?
4. Sind die für die RL-Arten der Gesamtliste erhaltenen Ergebnisse auf die RL-Waldarten übertragbar?

Methoden

Untersuchungsgebiet

Die Untersuchungen beziehen sich auf das niedersächsische Tiefland (Abb. 4.1), das sich in fünf naturräumliche Großregionen mit 41 naturräumlichen Regionen (NR) gliedert (MEYNEN & SCHMITHÜSEN, 1953–1962, modifiziert nach PILGRIM & FRANKE, 1993). Das Untersuchungsgebiet (UG) ist durch geringe Reliefunterschiede geprägt und von pleistozänen Ablagerungen dominiert (HEUNISCH et al., 2007). Von West nach Ost besteht ein Kontinentalitätsgradient, der sich in mittleren Jahresniederschlags- sowie Temperaturgradienten ausdrückt und sich auch auf die Bodenzusammensetzung auswirkt (NLFB, 1997).

Die Ems-Hunte-Geest und Dümmer-Geestniederung im Südwesten des UG ist in ihrem nördlichen Teil von ausgedehnten, vielfach von Flugsanden oder Sandlöss bedeckten Grundmoränenplatten geprägt. Ihr Südteil wird von Talsandflächen, großflächigen Mooren und kleineren Grundmoränenplatten dominiert, die z. T. von Endmoränenzügen überragt werden. Der Waldanteil liegt bei 17 %, geschützte Waldbereiche nehmen 1 % ein. Die Ostfriesisch-Oldenburgische Geest im Nordwesten liegt auf Grundmoränenplatten und wird intensiv landwirtschaftlich genutzt; Wälder haben einen Flächenanteil von 7 %. Das Weser-Aller-Flachland mit den Urstromtälern von Aller und Weser und den sich südlich anschließenden flachwelligen Moränenlandschaften liegt im Südosten des UG an der Grenze zum Hügel- und Bergland und hat mit 21 % den zweithöchsten Waldanteil im Tiefland. Die im Norden gelegene

Stader Geest ist durch Grundmoränen, Geesten sowie moorreiche Flussniederungen geprägt. Sie setzt sich aus kleinräumigen Acker-, Grünland-, Wald- und Mooregebieten zusammen. Wälder nehmen knapp 15 % der Region ein. In Lüneburger Heide und Wendland im östlichen Teil des UG überwiegen sandige Grund- und Endmoränen. Die Region wird von Äckern, Wäldern und den größten Sandheiden Niedersachsens geprägt und hat mit 39 % den höchsten Waldanteil der Großregionen des UG.

Gefäßpflanzenarten

Die Verbreitungsdaten der im UG als selten oder gefährdet eingestufteten Gefäßpflanzensippen (RL-Status der Tiefland-Region 1, 2, 3, G, R; GARVE, 2004) wurden der Datenbank des Niedersächsischen Pflanzenarten-Erfassungsprogramms entnommen (NLWKN 1982-2003, Stand 15.12.2008). Es gingen 490 Sippen (468 Arten, 14 Subspezies, 8 Aggregate) in die Analyse ein. Die Sippen wurden jeweils auf der höchsten in der Datenbank aufgeführten taxonomischen Ebene zusammengefasst. Die Analyse erfolgte auf Minutenfeld-Ebene (15 Minutenfelder pro Messtischblatt-Quadrant; insgesamt 15.729 Minutenfelder mit einer mittleren Größe von 207 ± 2 ha). Für die Teilanalyse der an den Wald gebundenen RL-Arten wurden aus der Gesamtliste diejenigen Sträucher und krautigen RL-Arten extrahiert, die im geschlossenen Wald sowie an Waldrändern und auf Waldverlichtungen vorkommen (Waldbindungskategorien 1.1, 1.2, SCHMIDT et al., 2011). Der Gruppe gehörten 60 Sippen (59 Arten, 1 Aggregat) an. Die Verbreitungsmuster der Arten wurden als Artenzahl pro Minutenfeld berechnet und in einem Geografischen Informationssystem (ESRI INC., 2007) visualisiert. Dabei wurden natürliche Grenzwerte mit jeweils fünf Klassen verwendet.

Schutzgebietssysteme

Die Ausdehnung und Lage der NSG und FFH-Gebiete wurden digitalen Daten entnommen (NLWKN, 2010a; NLWKN, 2010b, Stand: 31.12.2009). Für die Teilanalyse der an den Wald gebundenen RL-Arten wurden die digitalen Schutzgebietsdaten mit dem ATKIS-DLM-Layer Wald (LGN, Hannover) verschnitten. In die Analyse wurden zum einen durch NSG, zum anderen zusätzlich auch die durch FFH-Gebiete geschützten (Wald-)Flächen einbezogen und ihr Flächenanteil je Minutenfeld berechnet.

Effektivitäts-Analyse

Der Zusammenhang zwischen der Ausdehnung der Schutzgebiete (NSG bzw. NSG und FFH-Gebiete) bzw. der geschützten Waldflächen (durch NSG bzw. NSG und FFH-Gebiete geschützte Waldflächen) und der Anzahl von RL-Arten bzw. RLWaldarten pro Minutenfeld wurde mit Generalisierten Linearen Modellen mit Poisson-Verteilung untersucht. Es wurden einzelne Analysen für jede der 41 naturräumlichen Regionen berechnet. Die Schutzgebietsfläche je Minutenfeld war dabei

die erklärende, die Artenzahl die abhängige Variable. Die Datenanalyse erfolgte mit dem Statistikprogramm R (R DEVELOPMENT CORE TEAM, 2012).

Habitatbindung der RL-Waldarten

Für die weitere Interpretation wurde die Habitatbindung der RL-Waldarten gemäß FloraWeb-Datenbank (BfN, 2010; Haupt- und Schwerpunktorkommen) und GARVE (1994) ermittelt (bis zu drei Mehrfachnennungen je Art).

Ergebnisse

Verbreitung der RL-Arten

Im UG wurden insgesamt 490 RL-Arten nachgewiesen. Die absolute Artenzahl variierte zwischen den Großregionen stark. Während sie maximal bei 454 RLArten lag (Lüneburger Heide und Wendland), kamen im Minimum in der Ostfriesisch-Oldenburgischen Geest nur 255 Arten vor (Tab. 4.1). Die räumliche Verteilung von RL-Hotspots war deutlich inhomogen (Abb. 4.2). Maximal kamen pro Minutenfeld 86 Arten, im Mittel 5 (\pm 6) Arten vor. Weniger als 1 % der Minutenfelder wies mehr als 28 Arten auf, wobei die größte Dichte dieser Hotspots im Wendland lag (NR 860, 876). Rund 7 % der Minutenfelder waren sehr artenreich (15–28 Arten). Das westliche Tiefland war allgemein artenarm mit punktuell großflächigen Hotspots im Bentheimer Raum (NR 544, 580), im Emsgebiet (NR 586, 592, 604) und nordwestlich von Oldenburg (NR 602, 603). Im östlichen Tiefland war die Artendichte pro Minutenfeld tendenziell höher. Großflächige Hotspots lagen vor allem im Elbe-Weser-Dreieck (NR 631–633), in der Allerniederung (NR 626, 627), der Lüneburger Heide (NR 641, 642) und im Wendland. Die Teilmenge der an Waldhabitats gebundenen RL-Arten lag mit 60 Arten bei 12 %. Die Verbreitungsmuster der RL-Waldarten (Abb. 4.3) unterschieden sich stark von denen der Gesamtliste. Maximal kamen 22 Arten, im Mittel nur 0,5 (\pm 1,3) Arten pro Minutenfeld vor (Tab. 4.1). In 76 % der Minutenfelder konnte keine Art nachgewiesen werden. Die Hotspots mit mehr als sieben Arten waren nur punktuell ausgebildet und umfassten weniger als 1 % der Minutenfelder. Räume mit höheren Artendichten waren die Oldenburger Geest (NR 603), das Elbe-Weser-Dreieck (NR 633, 634), das Uelzener und Bevenser Becken (NR 643) sowie die Burgdorf-Peiner Geestplatten und das Ostbraunschweigische Flachland (NR 623, 624). Das Wendland wie auch die im äußersten Westen des UG gelegenen Gebiete, die bei Betrachtung der Gesamtliste die höchsten RL-Artenzahlen aufwiesen, waren vergleichsweise arm an RL-Waldarten.

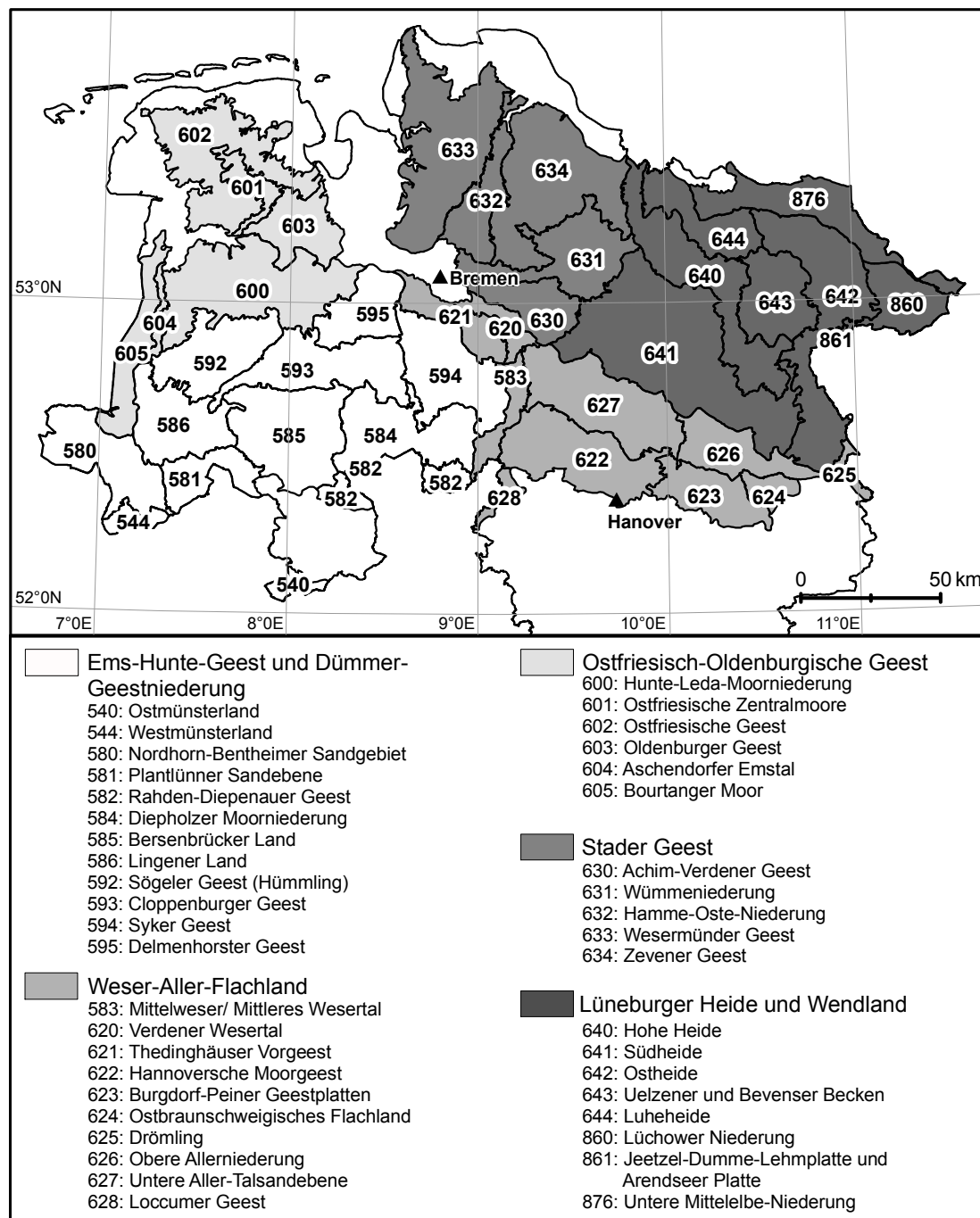


Abbildung 4.1: Das Untersuchungsgebiet des niedersächsischen Tieflandes mit seinen fünf naturräumlichen Großregionen und 41 naturräumlichen Haupteinheiten (MEYNE & SCHMITHÜSEN, 1953–1962, modifiziert nach PILGRIM & FRANKE, 1993). Bezugssystem: Deutsches Hauptdreiecksnetz (DHDN, Gauss-Krüger, Zone 3).

Tabelle 4.1: Gesamtfläche, absolute und prozentuale Flächenanteile von Schutzgebieten und Waldschutzgebieten (NSG und FFH-Gebiete) und Anzahl der RL-Arten bzw. RL-Waldarten in den fünf naturräumlichen Großregionen des niedersächsischen Tieflandes.

Großregion	Fläche [ha]	Waldfläche [ha]	Waldanteil [%]	NSG und FFH-Gebiete [ha]	Waldfläche in NSG und FFH-Gebieten [ha]	Anteil der Waldfläche in NSG und FFH-Gebieten an der Gesamt-Waldfläche [%]	RL-Arten	RL-Waldarten
Ems-Hunte-Geest und Dümmer-Geestniederung	863.759	143.640	16,6	47.572	9.674	6,7	336	39
Ostfriesisch-Oldenburgische Geest	457.808	31.455	6,9	27.265	2.885	9,2	255	26
Weser-Aller-Flachland	495.106	105.502	21,3	49.407	12.776	12,1	379	48
Stader Geest	553.736	81.363	14,7	40.194	8.190	10,1	327	37
Lüneburger Heide und Wendland	842.957	329.046	39	93.009	40.159	12,2	454	55
Tiefland Gesamt	3.213.366	691.006	21,5	257.447	73.684	10,7	490	60

Schutzgebietssysteme

Im UG waren 530 NSG ausgewiesen, die insgesamt 140.439 ha (4,4 %) der Gesamtfläche einnahmen. Die 240 FFH-Gebiete erfassten mit 212.962 ha 6,6 % der Tieflandfläche. Durch eine starke Überlappung der Schutzgebietskategorien (Abb. 4.4) summierte sich die insgesamt von NSG und FFH-Gebieten abgedeckte Fläche im UG auf 257.445 ha (8 % der Gesamtfläche; Tab. 4.1). Durch NSG geschützte Wälder nahmen 41 446 ha (1,3 % der Gesamtfläche), durch FFH-Gebiete geschützte Wälder 64.791 ha (2 %) ein. Der Anteil der insgesamt durch NSG und FFH-Gebiete geschützten Waldbereiche lag bei 2,3 %, der Anteil an der Waldfläche bei 10,7 % (Tab. 4.1).

Effektivität von Schutzgebieten für den Artenschutz

Die Vorkommen und Verbreitungsschwerpunkte der RL-Arten lagen zum überwiegenden Teil in Schutzgebieten (Tab. 4.2a). Die Effektivitätsanalyse zeigte für 85 % der NR einen positiven Zusammenhang zwischen NSG- und RL-Arten-Vorkommen, in den übrigen NR bestand kein Zusammenhang. In der Unteren Mittelbe-Niederung (NR 876), die eine außergewöhnlich hohe RL-Artenzahl und -dichte aufwies, lagen die Artvorkommen überwiegend außerhalb von NSG. Mit der Erweiterung der Schutzgebietskulisse um FFH-Gebiete (ca. 117.000 ha) erhöhte sich die Anzahl der NR mit Gebietsschutz auf 95 %. Dabei nahm z. B. die Schutzgebietsfläche in der Loccumer Geest von <1 % auf 18 % zu (Abb. 4.5a/b), im RL-Hotspot Untere Mittelbe-Niederung war sogar ein Anstieg von <1 % auf 24 % zu verzeichnen. Lediglich in zwei naturräumlichen Regionen, in denen keine nennenswerte Veränderung der Schutzgebietsfläche stattfand, wurde auch weiterhin keine effektive Abdeckung der Artvorkommen durch Schutzgebiete erreicht.

Die Hotspots der RL-Waldarten waren deutlich schlechter durch Schutzgebiete abgedeckt als die der RL-Arten der Gesamtliste (Tab. 4.2b). Lediglich in 61 % der NR lagen die Artvorkommen überwiegend in als NSG geschützten Wäldern. Durch die Ausweisung der FFH-Gebiete stieg der Anteil der NR mit einer guten Schutzgebietsabdeckung der Vorkommen der RL-Waldarten jedoch auf insgesamt 81 %. Markante Beispiele für diese Entwicklung sind die Oldenburger Geest (NR 603) und die Burgdorf-Peiner Geestplatten (NR 623). Hier sind die Minutenfelder mit besonders hohen Artendichten (>11 RL-Waldarten) nicht durch NSG geschützt, wurden jedoch in die FFH-Gebietskulisse einbezogen (Tab. 4.2b, Abb. 4.5c/d).

Gegenüber der Schutzgebietsabdeckung der Vorkommen der RL-Arten insgesamt waren die Vorkommen der RL-Waldarten damit zwar deutlich weniger stark durch Schutzgebiete gesichert, allerdings waren die verbleibenden NR, in denen kein Zusammenhang von RL-Waldartenverteilung und Schutzgebieten erkennbar war, generell relativ artenarm. Eine Ausnahme bildete lediglich die Rahden-Diepenauer Geest (NR 582), in der sowohl die RL-Arten insgesamt als auch die mit einer vergleichsweise hohen Zahl vorkommenden RL-Waldarten auch nach der Ausweisung von FFH-Gebieten nicht ausreichend durch Schutzgebiete gesichert waren.

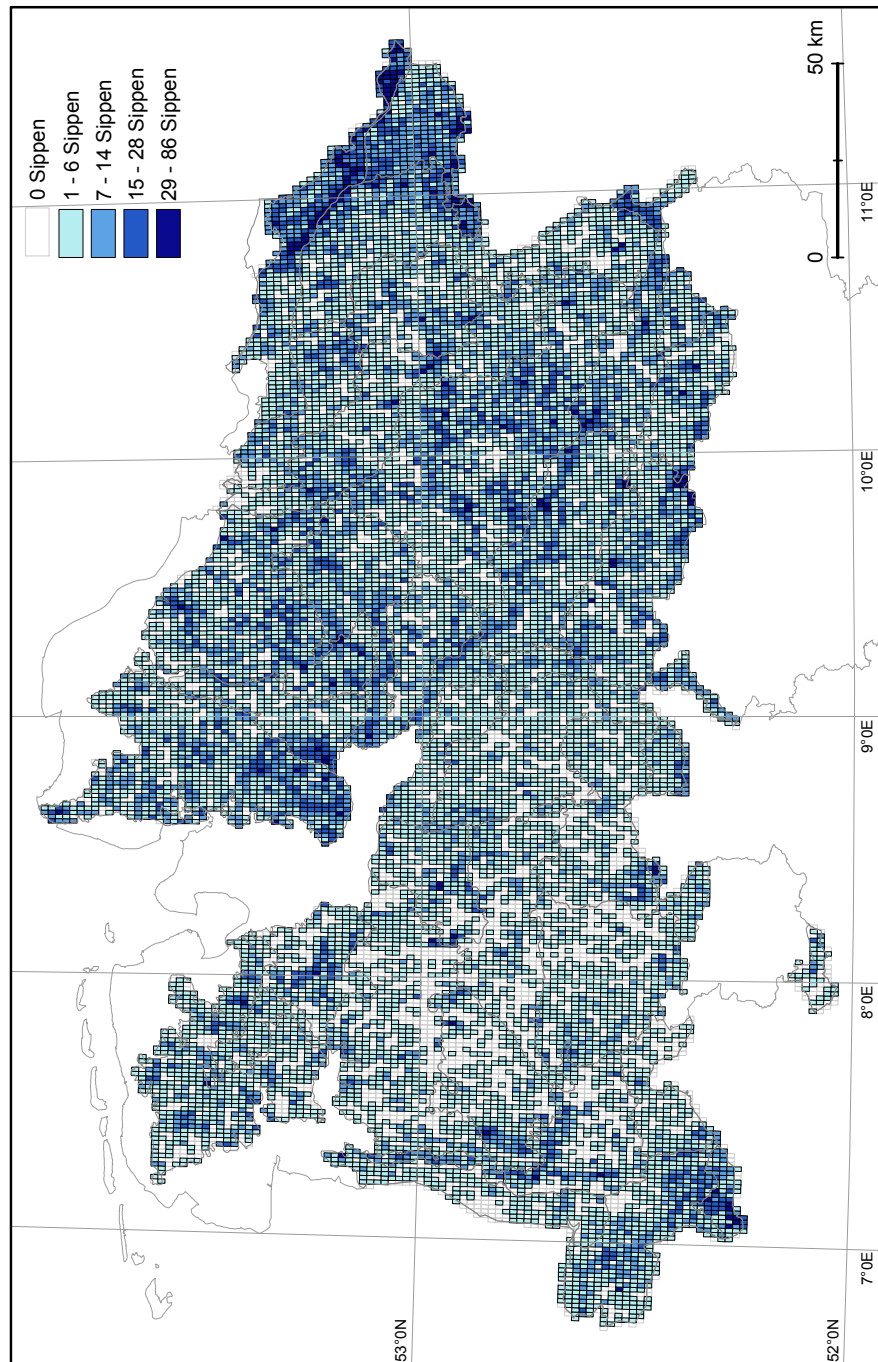


Abbildung 4.2: Anzahl der seltenen und gefährdeten Gefäßpflanzensippen pro Minutenfeld im niedersächsischen Tiefland. Insgesamt gingen 490 Sippen in die Berechnung ein.

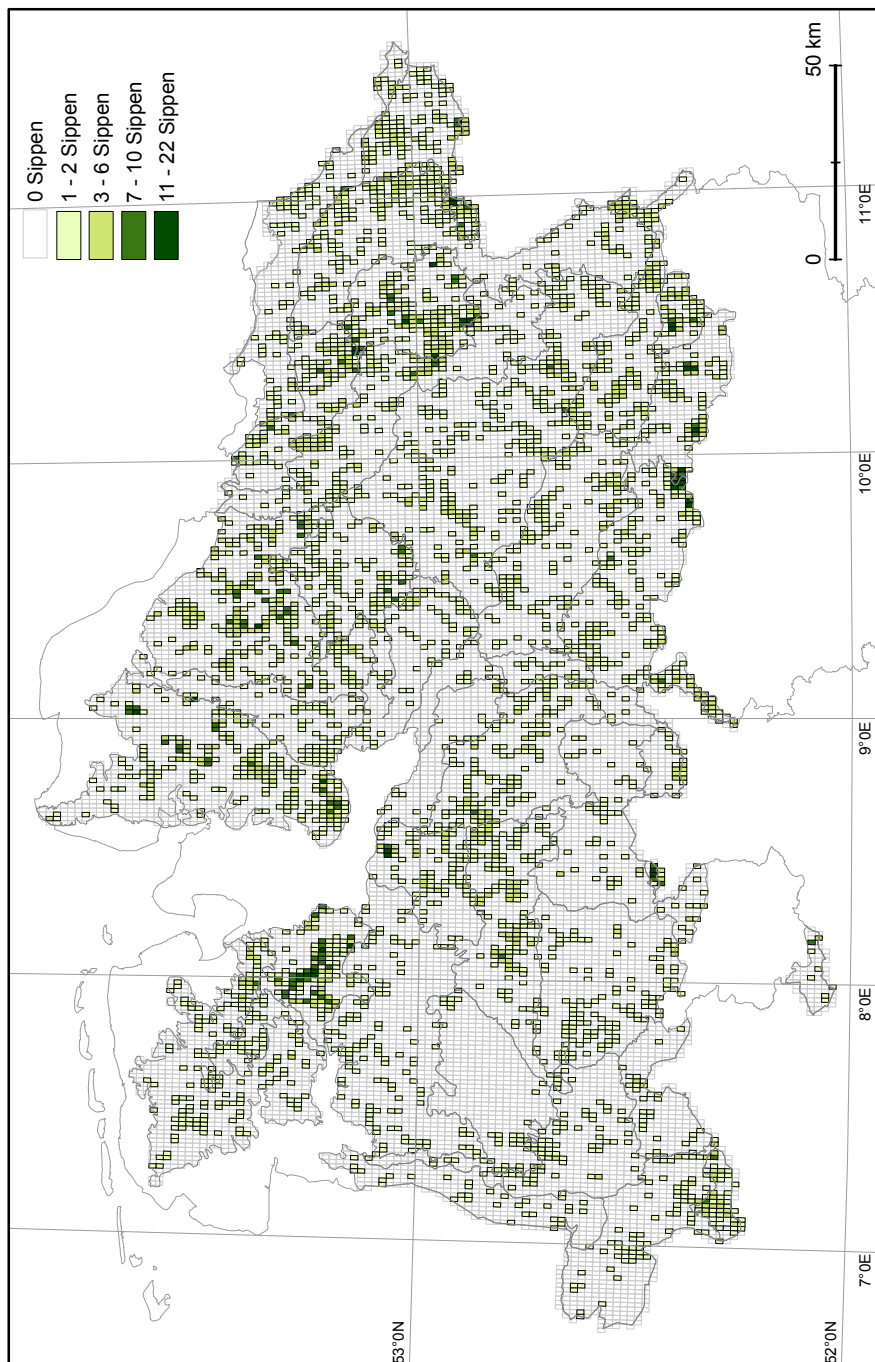


Abbildung 4.3: Anzahl der seltenen und gefährdeten Waldgefäßpflanzensippen pro Minutenfeld im niedersächsischen Tiefland. Insgesamt gingen 60 Sippen in die Berechnung ein.

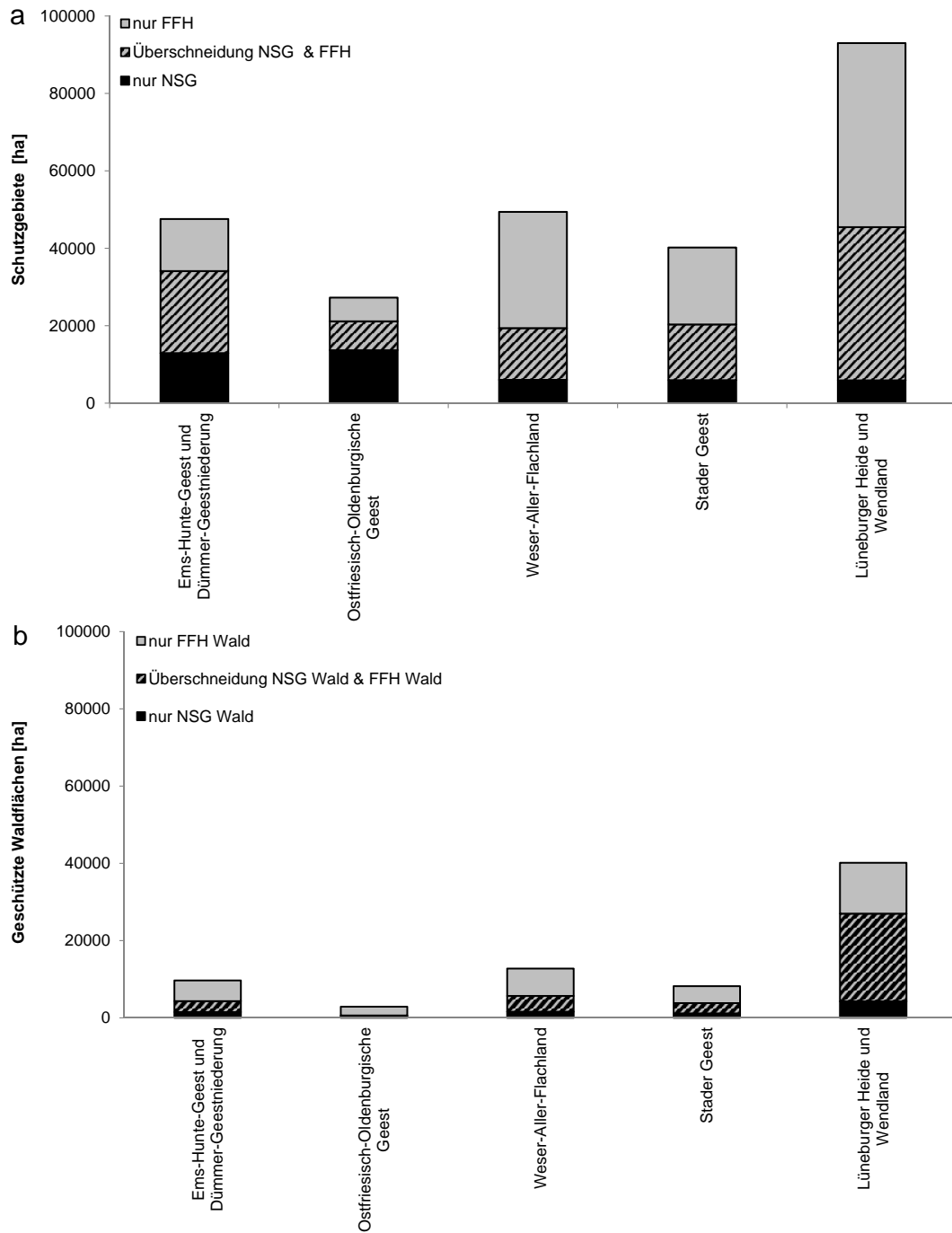


Abbildung 4.4: Schutzgebietsflächen (in Hektar) in den fünf Großregionen des niedersächsischen Tieflands unter Berücksichtigung der räumlichen Überlagerung von NSG und FFH-Gebieten. a) Gesamt Schutzgebietsfläche, b) Schutzgebiete in Wäldern.

Tabelle 4.2: Artenzahlen der RL-Arten bzw. RL-Waldarten, Ausdehnung der NSG bzw. der NSG und FFH-Gebiete und Ergebnisse der Analyse der Effektivität von Schutzgebieten für den Schutz von a) RL-Arten, b) RL-Waldarten. Effektivitätsanalyse: Ergebnisse der GLM-Analysen; n.s.: nicht signifikant, +/-: positiver/negativer Zusammenhang zwischen Schutzgebietsausdehnung und Artenreichtum, Signifikanzniveaus (* $p \leq 0,05$; ** $p \leq 0,01$; *** $p \leq 0,001$).

Großregion	NR	a) RL-Arten gesamt										b) RL-Waldarten				
		Artenzahl	NSG [ha]	NSG+FFH [ha]	NSG	Effektivitätsanalyse NSG+FFH	1.1	1.2	1.1+1.2	NSG Wald [ha]	FFH+NSG Wald [ha]	NSG Wald	Effektivitätsanalyse NSG Wald	NSG+FFH Wald		
Emse-Hunte-Geest und Dümmer-Geestniederung	540	75	21	21	n.s.	n.s.	10	0	10	18	18	n.s.	18	n.s.		
	544	148	687	1.918	+++	+++	18	2	20	111	111	n.s.	111	+++		
	580	175	1.754	3.104	+++	+++	13	0	13	614	614	+++	614	+++		
	581	111	96	264	n.s.	n.s.	12	0	12	6	6	n.s.	6	+++		
	582	149	156	165	n.s.	n.s.	22	1	23	9	9	n.s.	9	n.s.		
	584	134	18.929	20.53	+++	+++	10	2	12	460	460	+++	460	+++		
	585	171	1.293	3.47	+++	+++	21	0	21	393	393	+++	393	+++		
	586	148	2.017	6.52	+++	+++	10	1	11	302	302	+++	302	+++		
	592	118	5.739	6.355	+++	+++	9	0	9	455	455	+++	455	+++		
	593	122	673	999	+++	+++	14	1	15	283	283	+++	283	+++		
Ostriesisch-Oldenburgische Geest	594	144	921	1.842	+++	+++	15	1	16	526	526	+++	526	+++		
	595	154	1.831	2.384	+++	+++	20	1	21	1.168	1.168	+++	1.168	+++		
	600	156	11.66	12.173	+++	+++	8	0	8	334	334	+++	334	+++		
	601	179	2.828	2.879	+++	+++	3	0	3	20	20	n.s.	20	n.s.		
	602	179	1.885	3.564	+++	+++	21	0	21	180	180	+++	180	+++		
	603	131	398	1.242	+++	+++	18	1	19	12	12	n.s.	12	+++		
	604	75	635	3.669	+	+	4	0	4	4	4	n.s.	4	+++		
	605	86	3.73	3.738	+++	+++	2	0	2	58	58	n.s.	58	n.s.		
	620	121	1.017	1.459	+++	+++	10	2	12	39	39	n.s.	39	n.s.		
	621	112	66	1.34	+++	+++	3	1	4	3	3	n.s.	3	n.s.		
Weser-Aller-Flachland	622	261	242	388	n.s.	n.s.	5	1	6	108	108	+++	108	+++		
	623	233	1.767	3.252	+++	+++	32	4	36	407	407	+++	407	+++		
	624	104	149	305	+++	+++	31	3	34	208	208	n.s.	208	+++		
	625	109	1.792	3.992	+++	+++	22	2	24	128	128	+++	128	+++		
	626	228	5.367	9.628	+++	+++	12	1	13	947	947	+++	947	+++		
	627	229	3.14	14.401	+++	+++	20	1	21	3.132	3.132	+++	3.132	+++		
	628	88	13	1.816	n.s.	n.s.	20	3	23	693	693	+++	693	+++		
	630	135	985	1.693	+++	+++	13	2	15	5	5	n.s.	5	+++		
	631	195	5.674	11.399	+++	+++	13	0	13	305	305	n.s.	305	n.s.		
	632	131	2.415	5.431	+++	+++	22	0	22	1.237	1.237	n.s.	1.237	n.s.		
Stader Geest	633	235	8.164	11.655	+++	+++	16	0	16	127	127	+++	127	+++		
	634	208	3.098	10.016	+++	+++	28	1	29	1.067	1.067	+++	1.067	+++		
	640	202	17.494	19.326	+++	+++	25	2	27	1.104	1.104	+++	1.104	+++		
	641	250	18.494	33.262	+++	+++	33	4	37	11.832	11.832	+++	11.832	+++		
	642	305	1.273	3.426	+++	+++	28	3	31	8.909	8.909	+++	8.909	+++		
	643	198	1.533	3.401	+++	+++	31	4	35	880	880	n.s.	880	n.s.		
	644	196	1.525	2.527	+++	+++	31	2	33	1.028	1.028	+++	1.028	+++		
	860	285	3.761	8.179	+++	+++	33	4	33	853	853	+++	853	+++		
	861	120	779	827	+++	+++	16	0	16	168	168	+++	168	+++		
	876	273	618	22.061	-**	-**	16	6	22	447	447	+++	447	+++		
Summe		490	140.439	257.447		52	8	60	41.447	73.684		41.447	73.684			

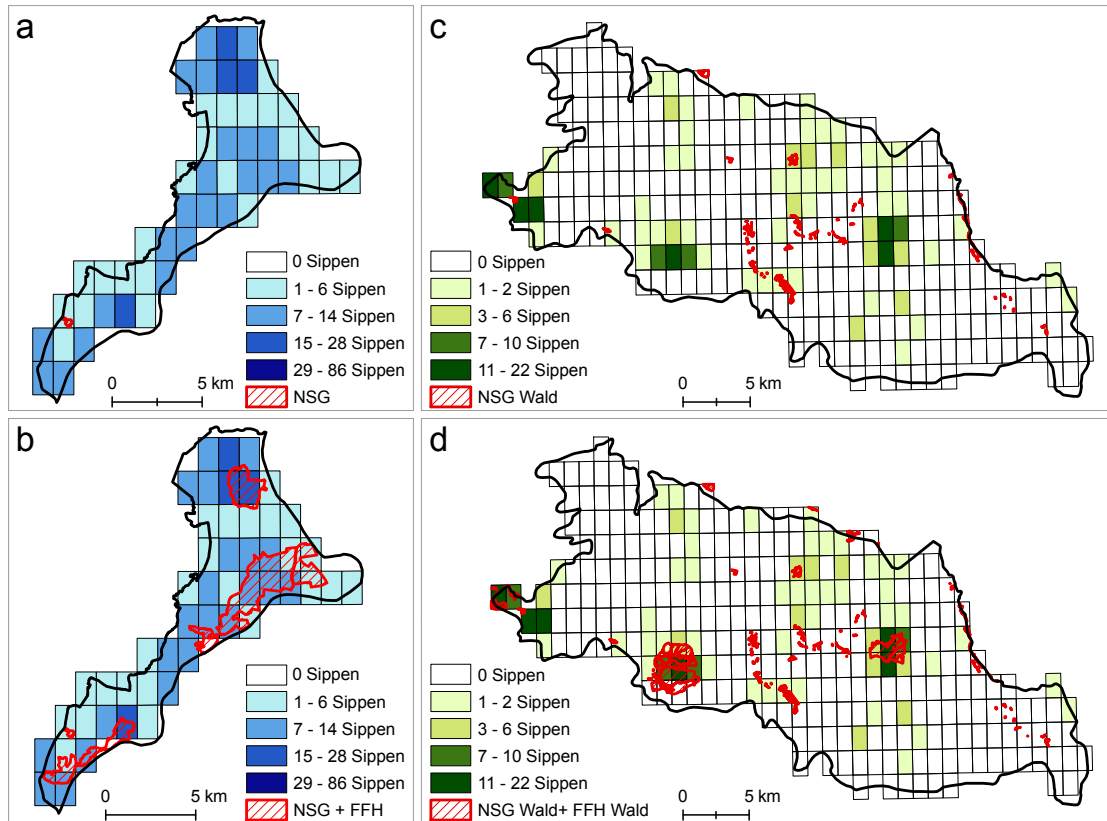


Abbildung 4.5: Beispiele für die Korrelation zwischen Schutzgebietsausdehnung und den Vorkommen der RL-Arten bzw. RL-Waldarten je Minutenfeld. In der Loccumer Geest (NR 628) überlagern sich die durch NSG geschützten Flächen und die Vorkommen der RL-Arten nicht (a), während sich unter Einbeziehung der FFH-Gebiete eine positive Korrelation zwischen Schutzgebietsausdehnung und den Vorkommen der RL-Arten zeigt (b). In den Burgdorf-Peiner-Geestplatten (NR 623) sind die Vorkommen der RL-Waldarten nicht durch NSG abgedeckt (c), während sich unter Einbeziehung der FFH-Gebiete eine positive Korrelation zwischen Schutzgebietsausdehnung und den Vorkommen der RL-Waldarten zeigt (d).

Diskussion

Die Verbreitung der seltenen und gefährdeten Gefäßpflanzenarten im niedersächsischen Tiefland zeigt ein differenziertes Bild (vgl. auch SCHMIEDEL et al., 2011). Es lassen sich Zentren der Artenvielfalt identifizieren, die im Sinne einer "Hotspots-Strategie" (MEYER et al., 2009) aus der landesweiten Perspektive als Naturschutz-Vorranggebiete angesehen werden müssen. Solche Hotspots finden sich vor allem im Wendland, im Westmünsterland und der Wesermünder Geest für RL-Arten bzw. der Oldenburger Geest für RL-Waldarten. Dagegen ist ein Großteil des westlichen Tieflands, wie etwa die Hunte-Leda-Moorniederung, das Bersenbrücker Land und das Bourtanger Moor, sehr artenarm.

Die Vorkommen der RL-Arten insgesamt sind nur in wenigen Landschaftsräumen des UG nicht durch Schutzgebiete gesichert (z. B. Ostmünsterland, Rahden-Diepenauer Geest). Die Vorkommen der RL-Waldarten sind zwar in einer geringeren Zahl der naturräumlichen Regionen effektiv durch NSG und FFH-Gebiete geschützt, allerdings ist die Zahl der RL-Waldarten in den Naturräumen mit nur geringfügiger Schutzgebietsabdeckung auch sehr klein. Gründe hierfür sind in diesen Regionen im geringen Anteil alter Waldstandorte und im insgesamt niedrigen Waldanteil zu suchen.

Im Allgemeinen verbessert sich die Schutzsituation für beide betrachteten Artengruppen bei Berücksichtigung von FFH-Gebieten. Dies zeigt, dass die erst in jüngerer Zeit ausgewiesenen FFH-Gebiete im niedersächsischen Tiefland künftig neben den NSG bei entsprechender Maßnahmenplanung eine wichtige Rolle für den Schutz seltener und gefährdeter Gefäßpflanzenarten spielen könnten. In diesem Zusammenhang sollte geprüft werden, ob der durch die Untersuchung bestätigte effektive Gebietsschutz auch in allen Fällen einen effektiven Artenschutz impliziert (vgl. RODRIGUES et al., 2004). Die Effektivität der Schutzgebiete für den Artenschutz ist in hohem Maße von den in den Schutzgebietsverordnungen bzw. Pflege- oder Managementplänen der Gebiete festgelegten Handlungsanweisungen abhängig. Insbesondere bei älteren NSG sind diese z. T. sehr unbestimmt formuliert (vgl. ENGEL et al., 2012) und für FFH-Gebiete derzeit noch in Arbeit. In vielen Fällen ist der Artenschutz zudem nicht vorrangiges Ziel bei der Schutzgebietsausweisung. Zukünftig sollten daher bei der Erarbeitung und Fortschreibung von Pflege- und Managementplänen in Schutzgebieten mit Hotspots zur dauerhaften Sicherung der bestehenden Populationen gezielt die Belange des Artenschutzes berücksichtigt werden.

Zu den Faktoren, die die Effektivität der Schutzgebiete einschränken können, zählen vor allem eine zu geringe Flächengröße der Gebiete bzw. ein hoher Grad an Isolierung: Sind die Schutzgebiete zu klein bzw. sind sie zu stark isoliert, können möglicherweise überlebensfähigen Populationen der schützenswerten Arten nicht erhalten werden (DEGUISE & KERR, 2006; JACKSON et al., 2009). Da ein Schutz größerer Flächen jedoch vielfach nicht umsetzbar ist, erscheint es in diesem Zusammenhang umso wichtiger, nicht allein auf die Wirkung von Schutzgebieten zu setzen, sondern im Rahmen einer "dualen Naturschutzstrategie" (JACKSON et al., 2009) auch Landschaftsbereiche außerhalb von Schutzgebieten so zu bewirtschaften, dass sie Populationen seltener und gefährdeter Arten nachhaltig als Lebensraum dienen können (DEGUISE & KERR, 2006; OTTE et al., 2008). In diesem Zusammenhang können beispielsweise die durch Eigenbindung ausgewiesenen Waldschutzgebiete im niedersächsischen Landeswald eine wichtige Funktion erfüllen (NIEDERSÄCHSISCHE LANDESFORSTEN, 2011).

Die für die Artengruppen voneinander abweichenden Ergebnisse zeigen, dass es sinnvoll ist, den Fokus nicht, wie in den bisher vorliegenden Studien zur Effektivität von Schutzgebieten (z. B. DEGUISE & KERR, 2006; JACKSON et al., 2009; VELLAK et al., 2009), lediglich auf die Gesamtheit der seltenen und gefährdeten, eine große

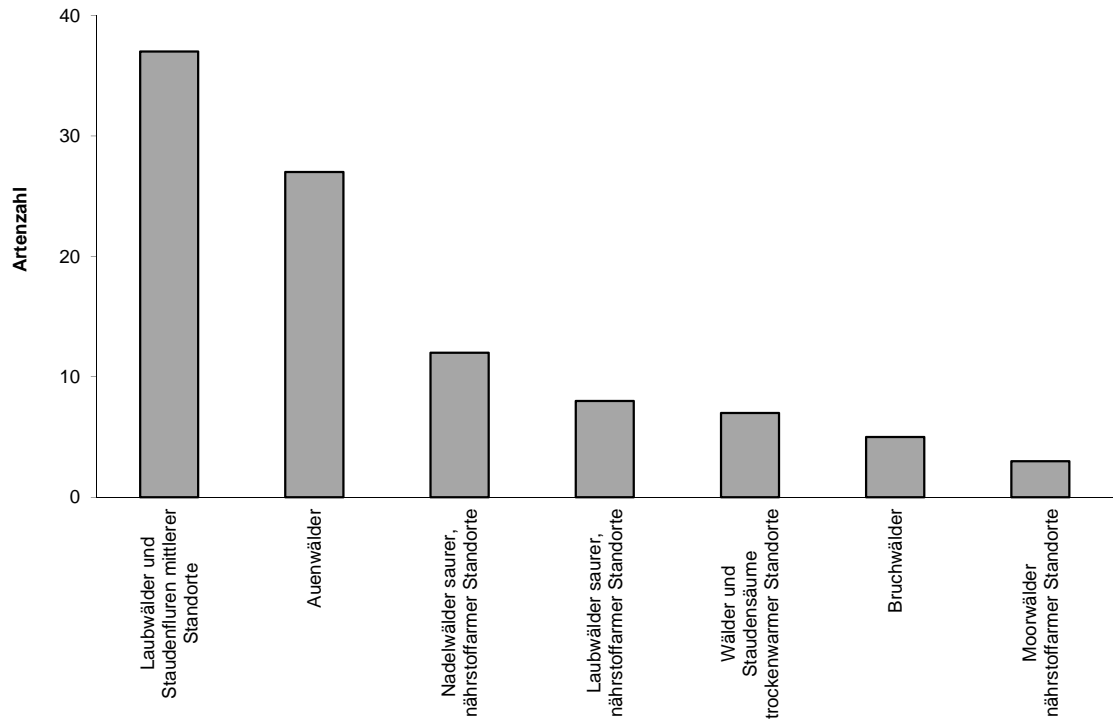


Abbildung 4.6: Habitatbindung der seltenen und gefährdeten Waldgefäßpflanzenarten des Tieflandes (bis zu drei Mehrfachzuordnungen pro Art).

standörtliche Amplitude repräsentierenden Gefäßpflanzenarten zu legen. Stattdessen kann die Analyse stärker differenziert und vergleichend auch solche Artengruppen einbezogen werden, die auf einen bestimmten Lebensraumtyp spezialisiert sind und somit eine geringere Standortamplitude aufweisen. Im vorliegenden Beispiel wurden hierfür die auf Wälder spezialisierten RL-Arten ausgewählt, da gerade Wäldern als der großflächig potenziell natürlichen Vegetation in Deutschland im Hinblick auf den Arten- wie auch den Prozessschutz eine große Bedeutung zukommt (BMU, 2007). Im niedersächsischen Tiefland zeigt ein Großteil der RL-Waldarten eine starke Habitatbindung an Laubwälder mittlerer Standorte (Waldmeister-Buchenwälder, Stieleichen-Hainbuchenwälder) und Auenwälder (Abb. 4.6). Von den 60 RL-Waldarten waren 43 an einen dieser beiden Habitattypen gebunden.

Die RL-Gefäßpflanzenarten können, obwohl sie nur einen Baustein des Gesamtökosystems darstellen, aufgrund ihrer Sensibilität gegenüber äußeren Einflüssen als gute Indikatoren für intakte Ökosysteme und die an diese gebundenen Arten und Lebensgemeinschaften angesehen werden.

Die fortlaufend durch das Pflanzenarten-Erfassungsprogramm erhobenen Verbreitungsdaten der RL-Arten sind eine sehr gute Grundlage für ein Monitoring der Effektivität der Schutzgebiete im UG. Wiederholende Untersuchungen in den folgen-

den Jahren werden zeigen, ob der in der vorliegenden Untersuchung belegte effektive Gebietsschutz langfristig auch zu einem effektiven Artenschutz führt.

Fazit für die Praxis

- Im niedersächsischen Tiefland liegen sowohl die Hotspots der Rote-Liste-Arten (RL-Arten) insgesamt als auch die der an Waldlebensräume gebundenen RL-Arten überwiegend in Schutzgebieten (NSG, FFH-Gebiete); die Ausweisung von FFH-Gebieten verbessert den Gebietsschutz für die RL-Arten in den meisten naturräumlichen Regionen erheblich. Wiederholungen der Untersuchungen werden zeigen müssen, ob der effektive Gebietsschutz auch zu einem effektiven Artenschutz führt.
- Bei der Erarbeitung und Fortschreibung von Pflege- und Managementplänen in Schutzgebieten mit Hotspots sollten künftig gezielt die Belange des Artenschutzes berücksichtigt werden, um die bestehenden Populationen von RL-Arten dauerhaft zu sichern.
- Die aufgezeigten Methoden der Hotspot-Analyse sollten stärker im Sinne einer "dualen Naturschutzstrategie" genutzt werden, um wichtige Populationen von RL-Arten auch außerhalb von Schutzgebieten durch nachhaltige Bewirtschaftungsformen zu erhalten.
- Rasterbasierte Verbreitungsdaten von RL-Arten sollten in regelmäßigen Zeitabständen aktualisiert werden, um sie als Grundlage eines Monitorings der Effektivität von Schutzgebieten für den Artenschutz einsetzen zu können.

Dank

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5

The influence of landscape fragmentation on richness patterns of vascular plants

Schmiedel, I. & Culmsee, H.: The influence of landscape fragmentation, expressed by the 'Effective Mesh Size Index', on regional patterns of vascular plant species richness in Lower Saxony, Germany. *Submitted to journal.*

Abstract

Landscape fragmentation has been identified as a major threat to biodiversity worldwide. Several landscape metrics have been developed to quantify the extent of fragmentation, of which the *Effective Mesh Size Index* (m_{eff}) is one of the most widely used. However, its relevance for biodiversity has been rarely tested. We analysed the explanatory power of m_{eff} for richness patterns of different groups of vascular plant species (all species, and species groups by residence and threat status) in Lower Saxony, Germany, by using a grid of 1386 landscape units. Since we assumed species richness to be influenced by abiotic conditions and spatial autocorrelation, we used variation partitioning to separate the effects of these variables from that of fragmentation. We tested five types of m_{eff} based on various fragmentation geometries of which two were identified as relevant for species richness.

We found that m_{eff} had a large effect on richness of neophytes and a slightly smaller effect on that of archaeophytes whilst the richness of native species was only slightly affected, and threatened species were not affected. All species groups, except threatened species, showed a negative correlation with m_{eff} , i. e. richness was highest in highly fragmented and lowest in less fragmented grid cells.

We conclude that m_{eff} is a meaningful tool to explain richness patterns of vascular plant diversity, if relevant fragmentation geometries are chosen. Our approach may therefore help future studies to determine correct fragmentation geometries to use with m_{eff} and may facilitate the unravelling of fragmentation impacts on the landscape-scale.

Keywords

Alien plant species, archaeophytes, native plant species, naturalisation, neophytes, PCNM, Red List, regional plant species richness patterns, variation partitioning, vascular plant diversity

Introduction

The term 'landscape fragmentation' describes both the process of the subdivision of a large continuous patch into smaller and more isolated fragments, as well as the state of a landscape (FORMAN, 1995). As soon as the effect of fragmentation on the biota is concerned, frequently also the term 'habitat fragmentation' is applied.

Besides general habitat loss and decline in habitat quality, fragmentation has been identified as a major threat to biodiversity worldwide (JAEGER, 2000; YOUNG & CLARKE, 2000; HONNAY et al., 2005; KUUSSAARI et al., 2009; KRAUSS et al., 2010). In Central Europe, habitat change accelerated in course of urbanisation and industrialisation in the late 19th and early 20th century (TSCHARNTKE et al., 2005; KRAUSE & CULMSEE, 2013) and has been reinforced by the advent of the common agricultural policy of the EU since the 1950s (STOATE et al., 2001; KRAUSE et al., 2011; LEUSCHNER et al., 2013). However, landscape development in Central Europe currently does not appear to be following a clear trend, and we observe complex and diverse landscapes as a result of multi-layered historical and current land use practices (VOS & MEEKES, 1999; CULMSEE, 2013). This, in turn, results in differences in the fragmentation intensity between neighbouring landscapes and in a more or less strong gradient of fragmentation on the regional and supra-regional scales.

The analysis of landscape fragmentation is an important aspect of landscape ecological research (HAILA, 2002). Many different indices for the quantification of landscape fragmentation have been introduced, such as the *Number of Undissected Areas*, *Bowens Landscape Dissection Index* (BOWEN & BURGESS, 1981), *Splitting Index* (JAEGER, 2000) and *Effective Mesh Size* (JAEGER, 2000). All of these indices provide a spatially differentiated assessment of the fragmentation of a region (JAEGER, 2000). By evaluating these four plus additional four indices, JAEGER (2000, p. 127) found that *Effective Mesh Size* (m_{eff}) is the most appropriate fragmentation measure due to its "mathematical characteristics and its intuitive interpretation". The index quantifies the probability that two randomly chosen points in a study area are connected (GIRVETZ et al., 2008) and has frequently been used to quantify landscape fragmentation, e. g. in Europe (EEA, 2011), Germany (JAEGER et al., 2001; ESSWEIN & SCHWARZ-VON RAUMER, 2004; WALZ, 2005; ESSWEIN & SCHWARZ-VON RAUMER, 2006), Italy (MOSER et al., 2007), Switzerland (JAEGER et al., 2008), the USA (GIRVETZ et al., 2008) and China (LI et al., 2010).

In Germany, m_{eff} is one of two indicators used by the government to quantify landscape fragmentation for environmental reporting (BMU, 2007). However, despite its frequent application and the fact that the ecological relevance of m_{eff} has been proposed by many researchers (JAEGER, 2000; JAEGER et al., 2008; GIRVETZ et al., 2008), its effect on biodiversity has been rarely analysed (but see LI et al., 2010). STRAND et al. (2007) suggested that studies investigating the direct effect of landscape fragmentation (expressed in terms of fragmentation metrics) on biodiversity should be undertaken, because indicators that quantify landscape fragmentation are more useful if they can be directly linked to possible impacts on species diversity and distributions.

A multitude of man-made landscape elements, including urban, industrial and traffic infrastructure, can cause landscape fragmentation, resulting in patchy landscapes with generally small remaining natural or semi-natural habitat fragments. Depending on the specific biological traits of the species, very different landscape elements, including (near-) natural ones, can be considered fragmentation elements (see GIRVETZ et al., 2008; EEA, 2011).

The fragmentation of habitats may result in increasing isolation of plant or animal subpopulations that inhabit the patches. Habitats colonised by different subpopulations of the same species are isolated if genetic exchange is constrained because the area between the patches is too large, or too impermeable, to be overcome by pollination or dispersal events (OOSTERMEIJER et al., 1994; POSCHLOD et al., 1997). The loss of genetic variation, in turn, may hamper the species' ability to respond to changing environmental conditions (BOUY et al., 2000) and the populations may thus become threatened with extinction. Numerous studies examined the effect of habitat fragmentation on plant and animal assemblages at the local and landscape scales (see for an overview DEBINSKI & HOLT, 2000; FAHRIG, 2003). Frequently, the size of and the isolation of or connectivity between patches were used as measure of fragmentation (e.g. HELM et al., 2006; COUSINS et al., 2007; KOLB, 2008; ZIMBRES et al., 2013), yet none of these studies applied the *Effective Mesh Size Index*.

The effects of landscape fragmentation on species richness may vary depending on the species groups considered (EWERS & DIDHAM, 2006; RODRIGUEZ-LOINAZ et al., 2012). Certain species, particularly those spreading along linear landscape elements or associated with man-made habitats, may benefit from landscape fragmentation (DEUTSCHEWITZ et al., 2003; HONNAY et al., 2003; DARK, 2004; NOBIS et al., 2009; BOTHAM et al., 2009). In contrast, threatened species may be adversely affected by fragmentation events (EWERS & DIDHAM, 2006). However, especially species that exhibit long life cycles and small population sizes show delayed extinctions following habitat loss and fragmentation (KUUSSAARI et al., 2009; KRAUSS et al., 2010), which at first can hamper the detection of these negative effects. This phenomenon is known as 'extinction debt' (TILMAN et al., 1994). However, the evidence for the existence of an extinction debt in vascular plants across Europe is ambiguous (reviewed in COUSINS, 2009).

The overall objective of our study was to investigate whether the frequently used *Effective Mesh Size Index* (m_{eff}) is a suitable measure to explain vascular plant species richness on the landscape scale. We used the state of Lower Saxony, Germany, as a model region as it hosts a high diversity of landscapes differing in habitat composition and structural complexity. Therefore, we expected a long gradient in fragmentation across the study area. Furthermore, the area is particularly suitable due to the availability of extensive data collected in a state-wide mapping program on the distribution of vascular plants with a resolution of c. 30 km² (NLWKN 1982-2003, GARVE, 2007). Thus, high quality data were available for a region that

is more or less homogeneous in climate and bio-geographical history, so that we considered the same regional species pool throughout the studied landscapes. We used different subsets of fragmentation geometries (FGs) and also different subsets of the regional plant species pool, dependent on the residence and threat status of the species, in order to investigate the underlying patterns in plant species distributions as responses to different fragmenting landscape elements.

Specifically, we hypothesised that:

1. Landscape fragmentation, expressed as *Effective Mesh Size* (m_{eff}), varies in space, which results in a gradient in the degree of fragmentation within the sample of landscape units used in this study.
2. The choice of fragmentation geometries that are used for the calculation of m_{eff} is of major importance for explaining plant species richness patterns on the landscape scale.
3. Plant species richness varies between landscape units along the fragmentation gradient, with the effect size differing between different groups of species dependent on their residence (natives, archaeophytes and neophytes) and threat status (threatened vs. non-threatened).

Methods

Study area

The study was carried out in Lower Saxony, Germany (c. 47,500 km², Fig. 5.1). As a result of the variety in relief and bedrock, colonisation history and current land use are diverse across north-western Germany (BEHRE, 2008). Lower Saxony harbours a relatively large number of different habitats, which include natural coastal and forest-dominated landscapes, structured and homogeneous agricultural landscapes, and densely populated areas (GHARADJEDAGHI et al., 2004).

The coastal area in the north is dominated by Holocene marine deposits, while the lowlands were formed during the Pleistocene. The landscape of the uplands is shaped by Mesozoic and Palaeozoic bedrocks. From the coast in the north to the uplands in the southeast, elevation ranges from 0–971 m a.s.l.. The climate gradually changes eastwards from oceanic (mean annual precipitation up to 900 mm) to suboceanic-subcontinental (c. 550 mm; NLfB, 1997).

Following the grid system of topographical maps, the study area was subdivided into a regular grid of 1386 cells sized c. 5.5 × 5.5 km² (30 km²; Fig. 5.1). In our analysis, we omitted the 353 border cells only partially occurring in the study area.

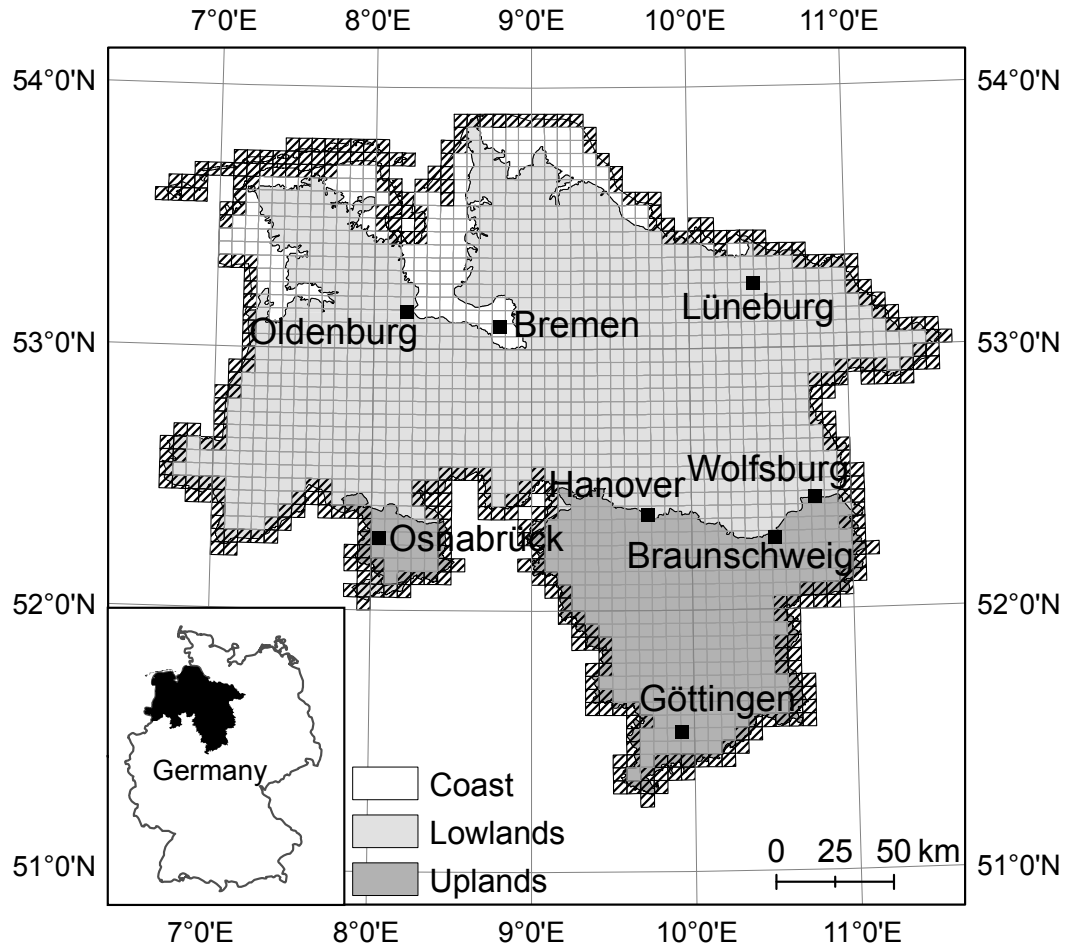


Figure 5.1: The state of Lower Saxony, Germany, with its three major landscape components: coast, lowlands and uplands. The 1386 grid cells (size c. $5.5 \times 5.5 \text{ km}^2$ or 30 km^2) display the basic mapping units of the vascular plant survey program and represent the landscape units for which landscape fragmentation was assessed. Hatched grid cells were not considered in this study, because they are not completely terrestrial or they are situated at the state's borders and thus information on plant occurrences were incomplete. Maps with Transverse Mercator projection, Germany Zone 3 ("Deutsches Hauptdreiecksnetz").

Plant species distribution data

Species occurrences were derived from the data base of the plant survey of the state of Lower Saxony (NLWKN, Hanover, Germany; period I from 1982 to 1992; period II from 1993 to 2003; period III from 2004 onwards). As the third fieldwork period is still underway, we used only data from 1982-2003 (GARVE, 2007; data base updated on 15 December 2008). The data base holds information about the presence and distribution of 2708 vascular plant species occurring in Lower Saxony (GARVE, 2007).

Table 5.1: Number of vascular plant species in Lower Saxony, Germany, grouped by their residence and threat status.

		Threat status			Sum of residence status
		Non-threatened	Threatened	Unspecified	
Residence status	Native species	703	482	112	1297
	Archaeophytes	103	46	0	149
	Neophytes	134	9	0	143
	Unspecified	4	1	0	5
Sum of threat status		944	538	112	1594

Species were assigned to different groups according to their residence and threat status (Table 5.1). In the designation of the residence status of the plant species we followed KLOTZ et al. (2002). Native species were distinguished from archaeophytes and neophytes. Within the non-native species, archaeophytes became established prior to and neophytes in the 16th century or later. In the designation of threat status we followed GARVE (2004), whereby threatened species (including the Red List categories 1, 2, 3, G and R) were distinguished from non-threatened species. Five species could not be assigned to a certain floristic status, while 112 species lacked information about their threat status (hereafter 'not specified' species). Subspecies were assigned to the species level and species to the respective aggregate found in the data base. In total, 1594 species remained that were considered in the analysis.

For each species group, species richness (based on presence/absence data) was calculated for each of the 1386 grid cells. Distribution patterns of the selected species groups were described in detail by SCHMIEDEL et al. (2011).

Quantification of landscape fragmentation

Landscape fragmentation analysis was conducted with the 1386 grid cells used by the plant species monitoring program (Fig. 5.1). Each grid cell was considered a landscape unit. Landscape elements were derived from a digital landscape model (DLM) of Lower Saxony, scale 1:25,000 (LGN, 2008), and summarised to eight major habitat types that represent suitable habitat formations for vascular plants: forest, grassland, cropland, water bodies, urban, industrial and traffic areas, and other habitats (e. g. islands, tree rows, bogs). Linear elements represented in the DLM as polylines (roads, rivers) were buffered by using either standard values (roads; FGSV, 1996) or information provided by the dataset itself (rivers) in order to convert them to polygons. Data were converted by using Hawth's Analysis Tools (BEYER, 2004) and combined with the other polygon features to a non-overlapping feature layer by using ArcGIS-ArcInfo 9.2 (ESRI INC., 2007).

Habitat types that were assumed to increase fragmentation of habitats suitable for different groups of vascular plant species were classified into five types of fragmentation geometries (FG1-FG5; Table 5.2). Starting with FG1, fragmenting landscape elements were successively added to the following FGs. FG1 included the most artificial land cover classes of traffic and urban-industrial areas. FG2 additionally included water bodies. Similar FGs have frequently been used in fragmentation studies and will thus allow for comparisons. The subsequent FG3-FG5 were built by adding further land cover classes (FG3: croplands, FG4: grasslands, FG5: other habitats) to the former FGs, thus setting a gradient of decreasing human impact (Fig. 5.2). FG5 included all landscape elements except forests which were rated as (semi-)natural landscape elements, because forests represent the potential natural vegetation of temperate Europe (BOHN & NEUHÄUSL, 2000/2003).

Landscape fragmentation was quantified by the *Effective Mesh Size Index* (m_{eff}). The index denotes the size of patches when the landscape is divided into S areas (each of the same size) with the same degree of landscape division as obtained for the observed cumulative area distribution (MCGARIGAL, 2014). Thus m_{eff} is an ecologically relevant metric that quantifies landscape fragmentation based on the probability that two randomly chosen points in a landscape unit are located in the same non-fragmented patch (JAEGER, 2000). The probability is multiplied by the total area of the landscape unit (in m^2) and finally gives the value of m_{eff} . The *Effective Mesh Size Index* was calculated following MCGARIGAL (2014) using the following formula:

$$m_{eff} = \frac{\sum_{j=1}^n a_{ij}^2}{A} \left(\frac{1}{10,000} \right) \quad (5.1)$$

where n is the number of patches, a_{ij} is the size of patch ij and A is the total area of the landscape. Values of m_{eff} are given in hectares (ha) by using the factor of 1:10,000 to convert m^2 into ha.

For the calculation of m_{eff} based on different FGs, subsequently called MeshFG1-MeshFG5 (Table 5.2), the FGs derived from the DLM vector data were transformed to raster format, and m_{eff} of the non-fragmented parts of the landscape was calculated for each grid cell using Fragstats 3.3 (MCGARIGAL et al., 2002). We applied the CUT procedure proposed by JAEGER (2000), and not the CBC procedure as proposed by MOSER et al. (2007), to obtain spatially explicit values for m_{eff} that could be correlated with the grid-based distribution data on plant species.

A Shapiro-Wilk normality test was performed on all m_{eff} variables to test them for normal distribution with the R software, version 2.15.2 (R DEVELOPMENT CORE TEAM, 2012). Furthermore, a Wilcoxon signed-rank test was used to test for significant differences between the m_{eff} types.

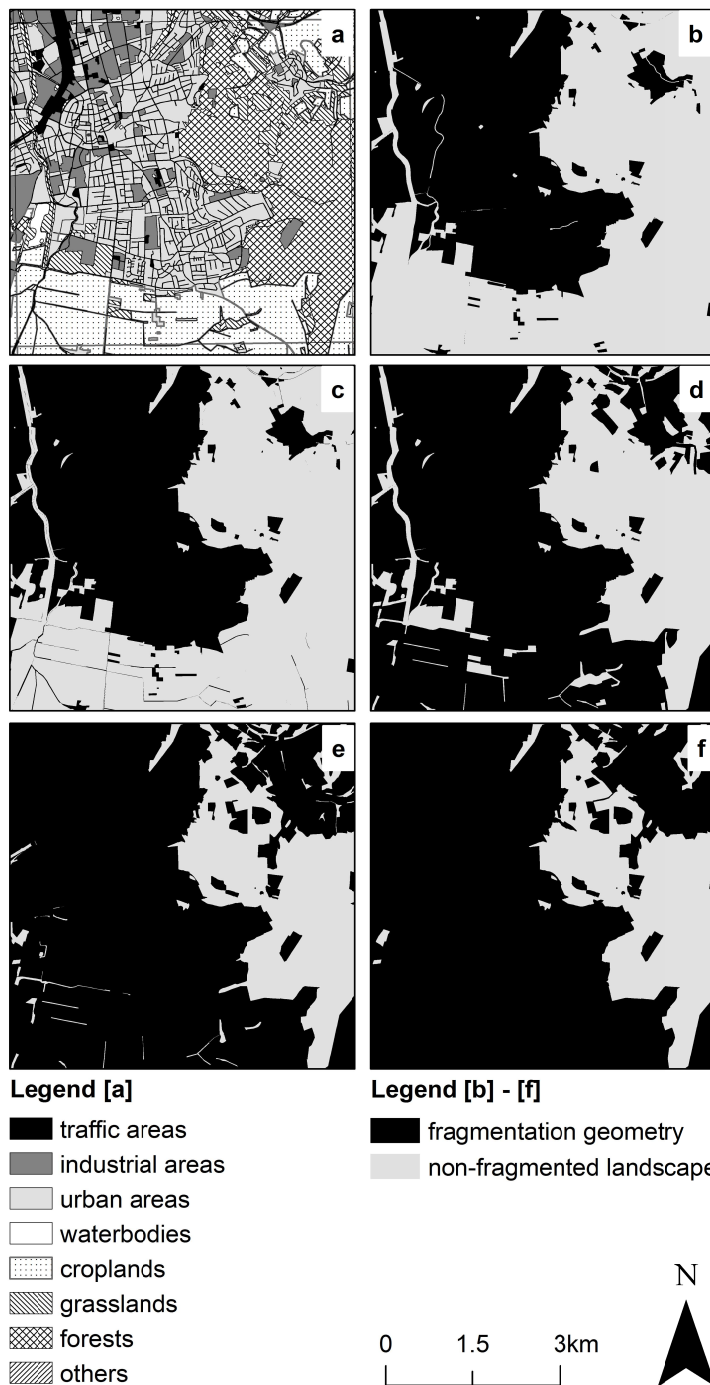


Figure 5.2: Example of one of 1386 landscape units in Lower Saxony, Germany (grid cell no. 44254, Göttingen, sized c. $5.5 \times 5.5 \text{ km}^2$ or 30 km^2). (a) The digital landscape model (ATKIS-DLM, scale 1:25,000; LGN, 2008) was translated to five different fragmentation geometries, FG1-5, (b) FG1 (traffic and urban-industrial areas), (c) FG2 (FG1 + water bodies), (d) FG3 (FG2 + croplands), (e) FG4 (FG3 + grassland), (f) FG5 (all landscape elements except forested areas).

Table 5.2: Landscape fragmentation variables (*Effective Mesh Size*, m_{eff} , based on different fragmentation geometries, FG1–FG5), spatial and environmental co-variables used in this study. n/a = not available. All m_{eff} types significantly differed from each other (Wilcoxon signed-rank test, $p \leq 0.001$).

Name	Definition	Unit	Source	Mean \pm SD	Range
Effective Mesh Size (m_{eff})					
MeshFG1	m_{eff} based on FG1: traffic and urban-industrial areas	ha	DLM ^b	1912 \pm 782	9–3196
MeshFG2	m_{eff} based on FG2: traffic and urban-industrial areas, water bodies	ha	DLM ^b	1430 \pm 819	0–3172
MeshFG3	m_{eff} based on FG3: traffic and urban-industrial areas, water bodies, croplands	ha	DLM ^b	412 \pm 565	0–3114
MeshFG4	m_{eff} based on FG4: traffic and urban-industrial areas, water bodies, croplands, grassland	ha	DLM ^b	195 \pm 451	0–3019
MeshFG5	m_{eff} based on FG5: all landscape elements except forested areas	ha	DLM ^b	152 \pm 381	0–2976
Spatial co-variables					
Ax1–	Significant spatial axes	n/a	n/a	n/a	n/a
Ax787					
Environmental co-variables					
ElevMn	Mean elevation	m a.s.l.	DEM ^c	65 \pm 84	-0.70–687.10
HeatlMn	Mean heat load index	n/a	DEM ^c	0.83 \pm 0.01	0.75–0.84
TempMn	Mean annual temperature	°C	DWD ^d	8.6 \pm 0.4	5.3–9.4
PrecipMn	Mean annual precipitation	mm	DWD ^d	743 \pm 90	562–1498
Snowdrop	Common Snowdrop (<i>Galanthus nivalis</i>) flowering	Day of year ^e	DWD ^d	56 \pm 6	29–85
Sun	Sunshine duration per year	h	DWD ^d	1450 \pm 54	1268–1615
SoilSHDI	Diversity of soil types (Shannon's Diversity Index, SHDI)	n/a	BÜK50 ^e	1.18 \pm 0.37	0.00–2.07
Calc1	Surface ratio of CaCO ₃ soil content: no CaCO ₃ content	ha	BÜK50 ^e	832 \pm 802	0–3205
Calc2	Surface ratio of CaCO ₃ soil content: medium CaCO ₃ content	ha	BÜK50 ^e	958 \pm 986	0–3208
Calc3	Surface ratio of CaCO ₃ soil content: high CaCO ₃ content	ha	BÜK50 ^e	528 \pm 6258	0–3065
Moist1	Surface ratio of soil moisture: dry	ha	BÜK50 ^e	11 \pm 81	0–1649
Moist2	Surface ratio of soil moisture: mesic	ha	BÜK50 ^e	738 \pm 699	0–3030
Moist3	Surface ratio of soil moisture: moist	ha	BÜK50 ^e	2932 \pm 482	0–3223
Moist4	Surface ratio of soil moisture: wet	ha	BÜK50 ^e	128 \pm 355	0–2967
Moist5	Surface ratio of soil moisture: influenced by backwater	ha	BÜK50 ^e	9 \pm 50	0–672

^a Day of year = day since the beginning of the year; ^b Digital landscape model of Lower Saxony (ATKIS-DLM, scale 1:25,000, LGN, 2008); ^c Digital elevation model for Lower Saxony (ATKIS-DEM, grid of 5 m \times 5 m resolution, LGN, 2008); ^d Climate data of the German Meteorological Service (grid of 1000 m \times 1000 m resolution, DWD 1961–1990); ^e Digital soil map of Lower Saxony (BÜK 50, scale 1:50,000, LBEG, 2003).

Accounting for environmental co-variables and spatial autocorrelation

We assumed that both landscape fragmentation and environmental factors influence species richness. Thus, to account for the explanatory value of the abiotic environment, supplementary environmental data were derived from various digital data sources (Table 5.2). Mean elevation was taken from the digital elevation model (DEM) of Lower Saxony (LGN, 2008). DEM data were further used to calculate mean heat load (McCune & Keon, 2002, eq. 3) per grid cell by using the ArcScript 'Heat load index' (Parks, 2004) in ArcView GIS 3.3 (ESRI Inc., 2002). Information on soil moisture, CaCO₃ content of soils and soil types were derived from the digital soil map of Lower Saxony (Bück 50, LBEG, 2003). For each grid cell, the surface ratio of five soil moisture and three CaCO₃ soil content classes were calculated. Based on the distribution and cover of 27 soil types, for each grid cell the Shannon Diversity Index (SHDI) was computed in Fragstats 3.3 after vector data were transformed to raster format. Information on climatic and phenological conditions (annual mean precipitation and temperature, start of snowdrop flowering) per grid cell were derived from digital data of the German Meteorological Service (DWD, 1961-1990) and the average of each variable was calculated per grid cell. The 15 environmental variables that were used in the analysis were uncorrelated (Spearman's rank correlation coefficient ($|\rho| \leq 0.7$)).

To account for spatial autocorrelation in the dataset, PCNM technique was applied using the following steps (cp. Borcard & Legendre, 2002; Borcard et al., 2004): (1) We constructed a matrix of Euclidean distances between the 1386 grid cells (landscape units) based on the x- and y-coordinates (centroids) of the cells. (2) We defined a threshold (smallest distance keeping all cells connected, i. e. 8036 m) below which the Euclidean distances were kept as measures and above which all distances were replaced by a value four times larger than the threshold (i. e. 32,144 m). (3) We computed the principal coordinates (correction of negative eigenvalues) of the modified matrix using PrCoord, version 1.0 (implemented in CANOCO 4.5, Braak & Šmilauer, 2002), and (4) we reduced the number of the remaining 787 spatial variables to those that were significant for the richness of the species groups (Monte Carlo Permutation test under full model, 99,999 permutations; Holm-correction, $p \leq 0.05$; Legendre & Legendre, 2012).

Statistical analysis

In order to determine the explanatory power of m_{eff} , the environmental factors and spatial axes for species richness patterns across the 1386 grid cells, variation partitioning was applied by using the 'varpart' function ('vegan' package, Oksanen et al., 2012) in R (cp. Fig. 5.3). Separate analyses were conducted for each species group (Table 5.1) in combination with each type of m_{eff} (Table 5.2). Species numbers of threatened species were square root transformed because they showed a

skewed distribution. Significance of all testable fractions was tested using 'rda' and 'anova.cca' functions in 'vegan'.

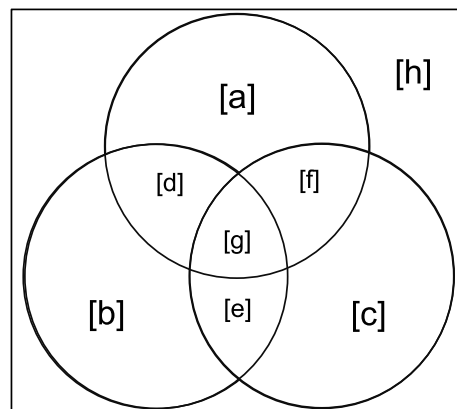
To determine the direction of the relationship of species richness and the m_{eff} types, Kendall's τ was calculated using R. This correlation analysis was performed for MeshFG1 and MeshFG2 only because variation partitioning showed that the other types of m_{eff} explained only a negligible proportion of the species richness of all groups.

Results

Variation in *Effective Mesh Size* (m_{eff}) based on different fragmentation geometries across landscapes

Values of m_{eff} calculated for the 1386 landscape units (grid cells) across the study area showed large variation both in range between landscapes and between fragmentation geometries (Table 5.2). All m_{eff} types (MeshFG1-MeshFG5) showed a large range from entirely fragmented ($m_{eff} = 0$ ha) to entirely non-fragmented ($m_{eff} \sim 3,000$ ha) landscape units, the latter corresponding to the total size of a landscape unit of c. 30 km². The mean value of m_{eff} was highest when only traffic and urban-industrial areas were used (MeshFG1, $m_{eff} = 1,912$ ha \pm 782 SD) indicating a relatively low fragmentation impact, and was lowest when all landscape elements with the exception of forests were used (MeshFG5, $m_{eff} = 152$ ha \pm 381 SD). From MeshFG1 to MeshFG5, mean m_{eff} gradually decreased when additional fragmentation geometries were successively included, whereas each of them had an additional impact as fragmenting landscape element against the previous (Wilcoxon signed-rank test, $p < 0.001$ for all pairs of MeshFG1-MeshFG5). None of the m_{eff} variables was normally distributed (Shapiro-Wilk normality test, $p \leq 0.05$). Values of MeshFG1 showed a slightly right-skewed distribution, i. e. larger values of m_{eff} were more frequent than small values (Fig. 5.4a), while values of m_{eff} for MeshFG2 were slightly left-skewed (Fig. 5.4b).

Figure 5.3: Fractions obtained from variation partitioning based on three variables (Venn diagram). Fractions [a] to [c] correspond to the pure effect of variables a, b and c, fractions [d] to [f] correspond to the variation explained by variables a and b, b and c, and a and c, respectively. Fraction [g] corresponds to the variation explained by all three variables together whereas [h] describes the unexplained variance (residuals).



The remaining m_{eff} types (Fig. 5.4c-e) all showed a strongly left-skewed distribution with the majority of landscape units exhibiting a mesh size <200 ha.

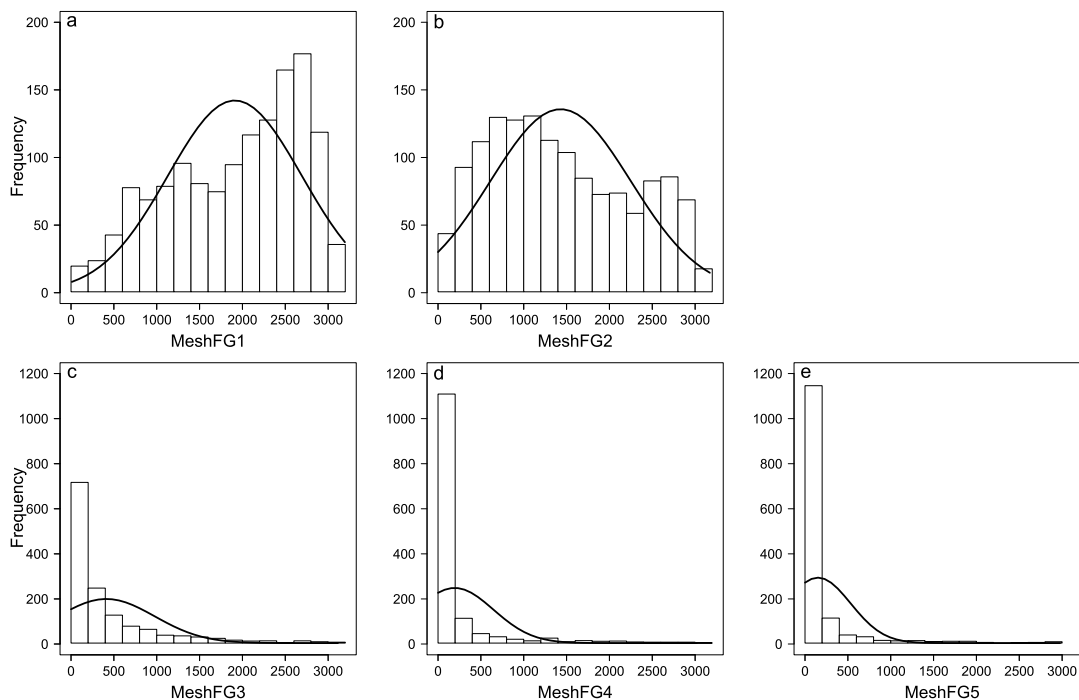


Figure 5.4: Frequency distribution of the five m_{eff} types (a-e, MeshFG1-MeshFG5) for the sample of 1386 landscape units (grid cells) in Lower Saxony, Germany. Maximum value of m_{eff} was c. 3,000 ha which corresponds to the size of a fully non-fragmented landscape unit of 30 km². None of the distributions were normally distributed (Shapiro-Wilk normality test, $p \leq 0.05$).

The explanatory power of *Effective Mesh Size* (m_{eff}), based on different fragmentation geometries, versus other environmental factors for species richness-patterns on the landscape-scale

Depending on fragmentation geometries (FGs) used for the calculation of m_{eff} and on the species group considered, the variation partitioning revealed large differences in the explanatory power of m_{eff} versus other environmental factors for vascular plant species richness patterns on the landscape scale (Table 5.3).

Accounting for the total of 1594 plant species, environmental variables explained 14% ($\pm 0SD$) and spatial variables 7% ($\pm 0SD$) of the variance in species richness (Table 5.3a). The response to landscape fragmentation largely varied in dependence on the mesh size variable (MeshFG1-MeshFG5). While m_{eff} types that used only traffic, urban-industrial areas and water bodies had a considerable influence of 5% (MeshFG1) and 6% (MeshFG2), the other m_{eff} variables, which additionally included

other man-made habitats such as agricultural areas and grasslands (MeshFG3 – MeshFG5), influenced species richness by <1%. Similar patterns were observed across all species groups. On average, 52% (± 3 SD) of variation in all species remained unexplained, and similar figures were found in the other species groups, with a range of 42% (± 4 SD) of unexplained variation in archaeophytes (Table 5.3c) to 61% (± 6 SD) in threatened species (Table 5.3f).

When distinguishing species groups according to the residence status, the general patterns observed in all species were more or less repeated in the partition of the largest group of 1297 native species, but with slightly lower explanatory power of the m_{eff} variables (Table 5.3b). Spatial distribution patterns of natives were also relatively clustered (explanation of the spatial fraction was $8\% \pm 0$ SD). However, the smaller groups of archaeophytes (149 sp.) and neophytes (143 sp.) showed different patterns. Archaeophytes were more strongly related to landscape fragmentation (MeshFG1 and MeshFG2: 7%) and especially to environmental conditions ($26\% \pm 2$ SD; Table 5.3c), indicating a relatively narrow environmental niche width of these species. Neophytes were most strongly associated with fragmented landscapes (Table 5.3d). M_{eff} explained a large proportion of species richness, namely 15% (MeshFG1) and 16% (MeshFG2), while the environmental factors were of lower importance ($10\% \pm 2$ SD). Spatial autocorrelation was generally low in both non-native groups (<2% of the total explained variation; Table 5.3c and d).

With respect to threat status, the large group of 944 non-threatened species followed the patterns observed in all species (Table 5.3e). In contrast, threatened species showed spatially distinguished incidence patterns (explained variation of spatial axes: $7\% \pm 0$ SD) and their occurrences were mostly determined by environmental variables ($19\% \pm 1$ SD), while they were not affected ($m_{eff} < 1\%$) by landscape fragmentation (Table 5.3f).

With the exception of the threatened species group that showed no specific trend, all species groups were negatively correlated with MeshFG1 and MeshFG2 (Table 5.4), which means that species richness increased with increasing landscape fragmentation (i. e. small m_{eff} values).

Discussion

Variation in landscape fragmentation based on m_{eff}

The values of m_{eff} obtained for the 1386 landscape units in our study varied between 0 and 30 km². Thus, we identified a large gradient from entirely fragmented to entirely non-fragmented landscape units. Because we applied the CUT procedure to calculate m_{eff} , maximum values of the index were restricted by the size of the study units. The latter also applies to other studies on m_{eff} (see Table 5 for an overview), which used differently sized reference units for the calculation of the index. Therefore, conclusions from comparisons of our results with values of m_{eff} obtained in these

Table 5.3: Results of variation partitioning (explained variation) for vascular plant species richness in Lower Saxony, Germany, in the groups of (a) all species, (b) natives, (c) archaeophytes, (d) neophytes, (e) non-threatened species, and (f) threatened species (numbers given in brackets), in relation to landscape fragmentation expressed as *Effective Mesh Size* (MeshFG1–MeshFG5) based on different sets of fragmentation geometries (FG1–FG5), spatial axes as measures for spatial autocorrelation and environmental factors (cp. Table 5.2). Significance was tested by Monte Carlo Permutation test (999 permutations) and refers to the partitions [a], [b] and [c] (cp. Fig. 5.3); significant results are given in bold letters; ***, $p \leq 0.001$; **, $p \leq 0.01$, *, $p \leq 0.05$.

	MeshFG1	MeshFG2	MeshFG3	MeshFG4	MeshFG5
(a) All species (1594 sp.)					
[a] M_{eff}	0.048 ***	0.061 ***	0.004 **	0.004 **	0.007 ***
[b] Spatial axes	0.067 ***	0.067 ***	0.073 ***	0.071 ***	0.074 ***
[c] Environmental variables	0.125 ***	0.145 ***	0.131 ***	0.139 ***	0.141 ***
Shared variation [d], [e], [f], [g]	0.264	0.242	0.252	0.245	0.241
[h] Residuals	0.497	0.484	0.541	0.540	0.537
(b) Natives (1297 sp.)					
[a] M_{eff}	0.024 ***	0.037 ***	0.001	0.002 *	0.005 ***
[b] Spatial axes	0.073 ***	0.073 ***	0.08 ***	0.077 ***	0.079 ***
[c] Environmental variables	0.121 ***	0.138 ***	0.123 ***	0.126 ***	0.129 ***
Shared variation [d], [e], [f], [g]	0.261	0.243	0.252	0.251	0.247
[h] Residuals	0.521	0.509	0.544	0.543	0.540
(c) Archaeophytes (149 sp.)					
[a] M_{eff}	0.067 ***	0.069 ***	0.013 ***	0.011 ***	0.011 ***
[b] Spatial axes	0.011 ***	0.016 ***	0.018 ***	0.018 ***	0.019 ***
[c] Environmental variables	0.245 ***	0.269 ***	0.247 ***	0.276 ***	0.274 ***
Shared variation [d], [e], [f], [g]	0.282	0.252	0.272	0.244	0.244
[h] Residuals	0.397	0.394	0.450	0.453	0.452
(d) Neophytes (143 sp.)					
[a] M_{eff}	0.154 ***	0.158 ***	0.010 ***	0.005 ***	0.002 *
[b] Spatial axes	0.010 ***	0.016 ***	0.021 ***	0.022 ***	0.023 ***
[c] Environmental variables	0.078 ***	0.078 ***	0.095 ***	0.115 ***	0.115 ***
Shared variation [d], [e], [f], [g]	0.152	0.147	0.124	0.103	0.103
[h] Residuals	0.606	0.602	0.750	0.755	0.758
(e) Non-threatened species (944 sp.)					
[a] M_{eff}	0.059 ***	0.069 ***	0.007 ***	0.006 ***	0.007 ***
[b] Spatial axes	0.063 ***	0.064 ***	0.071 ***	0.068 ***	0.072 ***
[c] Environmental variables	0.113 ***	0.135 ***	0.119 ***	0.13 ***	0.132 ***
Shared variation [d], [e], [f], [g]	0.275	0.252	0.262	0.252	0.248
[h] Residuals	0.490	0.479	0.542	0.543	0.541
(f) Threatened species (538 sp.)					
[a] M_{eff}	0.002 *	0.008 ***	0.004 **	0.000	0.001
[b] Spatial axes	0.066 ***	0.066 ***	0.068 ***	0.067 ***	0.066 ***
[c] Environmental variables	0.191 ***	0.197 ***	0.180 ***	0.174 ***	0.185 ***
Shared variation [d], [e], [f], [g]	0.090	0.084	0.100	0.107	0.096
[h] Residuals	0.650	0.644	0.648	0.652	0.651

Table 5.4: Correlation of the m_{eff} types MeshFG1 and MeshFG2, respectively, with species richness of six species groups (all species, groups after naturalization or threat status) assessed for Lower Saxony, Germany, using Kendall's τ ($p \leq 0.05$) as correlation coefficient.

m_{eff} type	Species group	Kendall's τ	z-score	p-value
MeshFG1 (max = 3,196 ha, min = 9.3 ha)	All species	-0.188	-10.49	< 2.2e-16
	Natives	-0.137	-7.62	2.61E-14
	Archaeophytes	-0.246	-13.62	< 2.2e-16
	Neophytes	-0.327	-18.00	< 2.2e-16
	Non-threatened species	-0.211	-11.75	< 2.2e-16
	Threatened species ^a	-0.015	-0.84	0.4016
MeshFG2 (max = 3,172 ha, min = 0 ha)	All taxa	-0.164	-9.15	< 2.2e-16
	Natives	-0.109	-6.04	1.55E-09
	Archaeophytes	-0.218	-12.07	< 2.2e-16
	Neophytes	-0.343	-18.84	< 2.2e-16
	Non-threatened species	-0.183	-10.19	< 2.2e-16
	Threatened species ^a	-0.011	-0.62	0.5364

^a Species richness of threatened species was square root transformed because it showed a skewed distribution.

studies have to be drawn with care. In contrast to the studies listed in Table 5, in our analysis, we applied units of equal size and thus the values of m_{eff} may be compared more easily. However, as we used relatively small landscape units, the actual degree of fragmentation may be overestimated. Nevertheless, compared to the results of other studies on m_{eff} in Germany (Table 5, no. 3-5), all of which used differently sized study units, our findings showed a similarly large variation. Values of m_{eff} obtained for Baden-Württemberg and Hesse ranged between 0 and c. 70 km² (JAEGER et al., 2001; ESSWEIN & SCHWARZ-VON RAUMER, 2004; Table 5, no. 3 and 5). In Bavaria, the maximum mesh size was c. 300 km², which was mainly a result of the high values of m_{eff} obtained for largely non-fragmented study units situated at the foothills of the Alps (ESSWEIN & SCHWARZ-VON RAUMER, 2006; Table 5, no. 4). The influence of largely non-fragmented mountainous areas also becomes obvious if values of m_{eff} for Switzerland are considered: When large alpine areas were included into the calculation procedure, maximum values of m_{eff} exceeded 1,500 km² (JAEGER et al., 2008; Table 5, no. 8). However, as soon as these uninhabitable areas were excluded from the computation process, the maximum values of m_{eff} dropped down to 300 to 700 km².

The European Environment Agency (EEA, 2011; Table 5, no. 2) compared the m_{eff} values of 28 EU countries and came to the conclusion that Germany, following the Benelux countries, was one of the most fragmented countries in Europe. Thus, the relatively small values of m_{eff} obtained for most of our studied landscape units seem to be representative for most parts of Germany. In contrast, studies from outside densely populated Europe showed much larger maximum values of m_{eff} (see Table 5, no. 1 and 9). For California, GIRVETZ et al. (2008) who used fragmentation geometries

similar to ours, and for China, LI et al. (2010) who calculated the *Effective Mesh Size* for countries and provinces, came up with maximum values of m_{eff} that exceeded these of our study a hundred- or even thousand-fold. However, even for these countries, the range of values of m_{eff} varied considerably: Whilst maximum values of m_{eff} in California reached c. 20,000 km² (Table 5, no. 9, B.1), the smallest values obtained were equal to zero. Similarly, values of m_{eff} for the 2,427 counties of China spanned a large range of c. 10 km² (the most densely populated areas) to c. 680,000 km² (the Tibetan plateau).

The response of species richness patterns to m_{eff} as a function of the choice of fragmentation geometries

Despite the fact that the FGs used in our analyses were nested, all types of m_{eff} (MeshFG1-MeshFG5) significantly differed from each other and thus did not contain any redundant information. Nevertheless, when relating the values of m_{eff} to species richness of the studied species groups, only MeshFG1 and MeshFG2 were found to be relevant. These m_{eff} types are based on FGs that include urban-industrial and traffic areas as well as water bodies and have been frequently used in studies on landscape fragmentation (cp. Table 5). Thus, even without examining the relationship of m_{eff} and species richness, many authors have chosen FGs that are meaningful for biodiversity. The explanatory power of MeshFG2 compared to that of MeshFG1 was slightly higher for all species groups. The inclusion of water bodies into the FGs used to calculate m_{eff} seems thus reasonable not only regarding animals, as proposed by GIRVETZ et al. (2008) and JAEGER et al. (2008), but also when distribution patterns of vascular plants are investigated. The remaining types of m_{eff} that included additional land cover classes supposed to have an added effect on the fragmentation of the landscape, did not explain (or only explained to a small extent) plant species richness patterns. We suppose that this was caused by the lack of a gradient of m_{eff} in MeshFG3 – MeshFG5; for the majority of grid cells very low values of m_{eff} were reported and the frequency distribution thus was extremely skewed to the left. This results from a country-wide intensive alteration of landscapes in Lower Saxony in the course of human land use that left only very few (almost) completely forested landscapes. Similar results could possibly be obtained for other intensively used landscapes all over Europe. However, if landscapes with a larger amount of (semi-)natural land cover classes are considered, the inclusion of agricultural areas (arable fields, managed grasslands) into m_{eff} calculation may reveal a more differentiated picture and thus may be more suitable to explain plant species richness. In contrast to our results, GIRVETZ et al. (2008) who investigated the effect of the inclusion of agricultural areas into the calculation of m_{eff} in California found these areas to only affect the degree of fragmentation in locations where extensive agricultural areas were present. In conclusion, the determination of relevant FGs is an important aspect when the relationship of landscape fragmentation and biodiversity is analysed.

Table 5.5: Overview on studies using the *Effective Mesh Size Index* (m_{eff}) to quantify landscape fragmentation, their coverage, study units and the fragmentation geometries (FGs) as well as potential response variables used in the particular studies. The range of m_{eff} is given for each FG and study unit, respectively. n/a = not available. Unless otherwise specified, m_{eff} is reported for CUT procedure.

No	Coverage (total area)	Study units	Size of study units (ranges, km ²)	Fragmentation geometries (FGs) used	Range of m_{eff} (in km ²)	Response variables	Reference
1	China (c. 9,500,000 km ²)	A: 34 provinces, B: 2,427 counties	A: n/a B: n/a	National, provincial and county roads, railway lines, urban areas	A: 29.9–304,491 B: 10.7–676,623 ^a	Endangered plants, mammals	LI et al. (2010) ^d
2	European Union, 28 countries (c. 4,400,000 km ²)	A: 28 EU countries, B: 581 NUTS-X regions, C: 1 km ² grid units within the 28 countries	A: c. 300–544,000 B: n/a C: 1	Motorways, other roads up to class 4, railway lines, urban areas, mountains, lakes, major rivers	A: 7.40–2,525.04 B: 0.03–6,177.91 C: n/a	None	EEA (2011) ^d
3	Germany, Baden-Württemberg (c. 36,000 km ²)	A: 44 administrative districts, B: 66 natural geographic units	A: 97.9–1,860.5 B: 13.9–1,989.7	1: Urban areas, water bodies, railway lines, major roads 2: FG1 + municipal roads	A.1: 2.09–46.73 B.1: 1.20–71.86 A.2: 1.63–30.26 B.2: 0.78–566.14	None	JAEGER et al. (2001)
4	Germany, Bavaria (c. 70,500 km ²)	A: 7 administrative districts, B: 18 regions, C: 16 nature parks, D: 96 natural geographic units	A: 7,231.1–17,530.1 B: 1,478.5–5,689.7 C: 229.9–2,966.2 D: 13.5–4,494.6	1: All roads excluding municipal roads, multi-tracked railway lines, channels, urban areas 2: Roads used by $\geq 1,000$ vehicles per day, heavily used railway lines, urban areas, airports, larger channels 3: Similar to FG 2, roads buffered according to traffic density	A.1: 16.37–48.43 B.1: 15.02–108.42 C.1: 16.24–40.61 D.1: 0.59–286.71 A.2: 45.75–71.85 B.2: 23.13–112.13 C.2: 31.87–153.15 D.2: 0.60–287.01 A.3: 36.40–65.70 B.3: 14.50–92.49 C.3: 27.15–132.69 D.3: 0.09–286.38	None	ESSWEIN & SCHWARZ-VON RAUMER (2006)
5	Germany, Hesse (c. 21,000 km ²)	A: 3 administrative districts, B: 26 administrative districts, C: 59 natural geographic units	A: 53,804.8–82,909.9 B: 44.8–1,848.9 C: 0.5–1,553.2	1: All roads excluding municipal roads, multi-tracked railway lines, channels, urban areas 2: Similar to FG 1, roads buffered according to traffic density	A.1: 14.87–19.42 B.1: 1.94–27.68 C.1: 0.44–67.06 A.2: 11.40–15.80 B.2: 0.11–20.11 C.2: 0.00–45.38	None	ESSWEIN & SCHWARZ-VON RAUMER (2004)

Continued on next page

Table 5.5 – Continued from previous page

No	Coverage (total area)	Study units	Size of study units (ranges, km ²)	Fragmentation geometries (FGs) used	Range of m_{eff} (in km ²)	Response variables	Reference
6	Germany, Lower Saxony (c. 47,500 km ²)	1,386 grid cells of c. 5.5 × 5.5 km ²	c. 30	1: Traffic and urban-industrial areas 2: Traffic and urban-industrial areas, water bodies, croplands 3: Traffic and urban-industrial areas, water bodies, croplands, grassland 4: Traffic and urban-industrial areas, water bodies, croplands, grassland 5: All landscape elements except forested areas	0.09–31.96 0.00–31.72 0.00–31.14 0.00–30.19 0.00–29.76	Six groups of vascular plant species	This study
7	Italy, South Tyrol (c. 7,500 km ²)	Municipality boundaries	1.6–302.3	Areas of urban development and transportation infrastructure (roads, railway lines)	0.50–208.70	None	Moser et al. (2007)
8	Switzerland (c. 41,000 km ²)	A: 5 ecoregions, B: 26 cantons, C: 6 BLN-areas	A: n/a B: n/a C: n/a	1: Motorways, highways (class 1–3), railway lines, urban areas 2: Motorways, highways (class 1–2), railway lines, urban areas, water bodies, mountains (≥2,100m) 3: FG 1 + water bodies and mountains (≥2,100m) 4: equals FG 1 but inhabitable parts of landscape omitted	A.1: 19.96–1,310.24 B.1: 0.54–1,850.75 C.1: 27.58–1,544.30 A.2: 21.47–430.58 B.2: 0.61–681.82 C.2: n/a A.3: 9.86–293.30 B.3: 0.52–485.52 C.3: n/a A.4: 10.78–381.72 B.4: 0.54–635.14 C.4: n/a	None	JAEGGER et al. (2008) ^d
9	USA, California (c. 450,000 km ²)	A: 2 nested administrative planning unit boundaries, B: 6 nested spatial scales of watersheds	A: 118–520,911 B: n/a	1: Highways, major roads, railway lines, urbanized areas 2: FG 1 + minor roads 3: FG 2 + agricultural areas 4: FG 3 + lakes, major rivers, alpine areas above 3,000 m	A.1: 0.04–12,092.00 ^b B.1: 0.00–20,885.00 ^b A.2: 0.02–5,064.00 ^b B.2: 0.00–14,900.00 ^b A.3: 0.02–5,058.00 ^b B.3: 0.00–14,898.00 ^b A.4: 0.01–2,615.00 ^b B.4: 0.00–10,175.00 ^b	None	GRIEVEZ et al. (2008) ^d

^a Average of m_{eff} for the 10 least fragmented counties; ^b 58 counties of California, USA; ^c 6,994 planning watersheds of California, USA; ^d CBC procedure (cp. Moser et al. 2007).

Differentiating the effect of m_{eff} on species richness for selected plant groups

The explanatory power of m_{eff} compared to that of abiotic and spatial variables differed considerably among species groups. Surprisingly, species richness and landscape fragmentation were positively correlated for all species groups, with the exception of threatened species. Thus, species richness was highest in highly fragmented landscapes and was lowest in little fragmented landscapes.

For richness patterns of neophytes m_{eff} was an exceptionally important factor. That is not surprising, because alien species are known to be linked to urban habitats and linear landscape elements and are associated with human activities (e. g. PYŠEK, 1998; ROY et al., 1999; DEUTSCHEWITZ et al., 2003; HONNAY et al., 2003; KÜHN et al., 2003; LIPPE & KOWARIK, 2008). Archaeophytes, in contrast, are frequently found in rural environments (PYŠEK, 1998; DEUTSCHEWITZ et al., 2003), particularly in loess landscapes (KÜHN et al., 2003) with a long colonization history (KÜSTER, 1999). Today, most of these areas in our study area are intensively used and highly fragmented agricultural and/or industrial landscapes. The relation of archaeophyte richness with loess landscapes and major rivers in Lower Saxony was confirmed by SCHMIEDEL et al. (2011). As suggested by the high explanatory value of the environmental variables, areas of high archaeophyte richness are related to environmental conditions rather than to landscape fragmentation.

Only a very small proportion of native species richness was explained by m_{eff} , while environmental variables were more important. Even if weak, a negative relationship between m_{eff} and species richness was observed for this species group as well. This finding may be attributed to the fact that urban areas are commonly situated in naturally diverse ecoregions as shown by KÜHN et al. (2004), who found that geological diversity is associated with areas of urbanisation in Germany. Furthermore, niche availability in urban areas is known to be rather high (DEUTSCHEWITZ et al., 2003; HONNAY et al., 2003; KÜHN et al., 2004; WANIA et al., 2006; LIPPE & KOWARIK, 2008). Possible adverse effects of fragmentation could therefore be masked.

Surprisingly, we found no effect of m_{eff} on threatened plant species richness patterns. Environmental conditions were better predictors of threatened plant species richness than was m_{eff} . Similar to our results, HONNAY et al. (2003), who studied the effect of different landscape indices as predictors for regional plant species richness, found the effect of landscape fragmentation on the richness of threatened species to be compensated by high landscape diversity. However, it seems that threatened species' occurrences are independent of the overall degree of landscape fragmentation (at least as long as it remains above a certain threshold). Threatened species are usually closely linked to ecologically valuable, often threatened habitat types. As discussed for Natura 2000 forest habitat types, characteristic forest species are limited in their dispersal ability and mostly linked to forests with long habitat continuity (CULMSEE et al., 2014). At first, fragmentation may have only little effect on remnant

populations as long as habitats do not vanish completely, yet the consequences of an extinction debt might be paid in the future (KUUSSAARI et al., 2009). However, the observed result may be an effect of the study scale being too coarse to reveal the relationship between landscape fragmentation and richness of threatened species (JACKSON & FAHRIG, 2012). Further studies on different spatial scales are needed to provide evidence on the latter.

Conclusions

The assessment of landscape fragmentation and approaches for linking fragmented habitats by corridors are becoming increasingly important in nature conservation policy in Germany, as well as the rest of Europe (BfN, 2014; EUROPEAN COMMISSION, 2011; BUNDESKABINETT, 2012). The EEA (2011, p. 58) even identified the analysis of the relationship between the levels of landscape fragmentation and biodiversity as "one of the most important areas for future research". In this context, measures of fragmentation are increasingly used to assess the status of a landscape (STRAND et al., 2007; BMU, 2010). The *Effective Mesh Size* m_{eff} is one of the most frequently applied indices in studies on landscape fragmentation. While many of these studies have pointed out the ecological relevance of fragmentation in general and m_{eff} in particular (JAEGER, 2000; JAEGER et al., 2008; GIRVETZ et al., 2008; LI et al., 2010), to our knowledge, until now no landscape-scale study has evaluated the impact of m_{eff} on species richness. By addressing this issue, our study has contributed to increasing the understanding of this relationship. We conclude that, as long as relevant fragmentation geometries for the landscapes under consideration are chosen, m_{eff} can act as a meaningful measure to explain vascular plant species richness patterns. Out of the five types of m_{eff} used in our study, two were found to be relevant for species richness, both having been used in previous studies (JAEGER et al., 2001; GIRVETZ et al., 2008; JAEGER et al., 2008). We further conclude that it is important to consider not only all vascular plant species, but also several smaller species groups. Species included in these groups should differ in their residence or threat status or in other characteristics in order to obtain more precise results on habitat requirements that facilitate interpretations.

Because fragmentation is a landscape-scale process, its relationship with biodiversity should accordingly be studied at this scale (MCGARIGAL & CUSHMAN, 2002; FAHRIG, 2003). However, when evaluating 100 and 135 fragmentation studies, respectively, MCGARIGAL & CUSHMAN (2002) and FAHRIG (2003) recognised that about 50 % of all studies were actually conducted at the patch scale. This is obviously related to the practical difficulties arising from large-scale experiments. Thus, to be able to study the effects of landscape fragmentation on biodiversity, it is important to develop new methods for their assessment. Our approach avoids such difficulties by using readily available high-resolution data on land cover and the distribution of vas-

cular plant species. As such data are becoming increasingly available in more and more countries (CULMSEE et al., 2014) and because computational power is rapidly increasing, our approach is easily extendable to other – even larger – study areas.

Further studies are needed to improve our understanding of the relationship of species richness and landscape fragmentation. The transferability of our results to other spatial scales should be tested, as well as the applicability of the method to less human-dominated landscapes. Presumably, effects of landscape fragmentation on species in highly transformed Central European landscapes differ from those in more natural landscapes. Thus, further studies are encouraged to bridge the gap between the assessment of landscape fragmentation, which is methodologically already advanced, and its relevance to species diversity, which has been rarely tested in the past.

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6 | Plant species richness along a gradient of landscape modification

Schmiedel, I.; Bergmeier, E. & Culmsee, H.: Plant species richness along a gradient of landscape modification in Lower Saxony, Germany. *Submitted to journal.*

Abstract

Several concepts of landscape modification, landscape naturalness and hemeroby have been put forward to quantify the extent of human activities and impact on landscapes. However, despite numerous studies on the human influence on landscapes, the relationship between these concepts and biodiversity has rarely been tested. On a grid of 1386 lat/long cells (landscape units, c. 30 km²) we used simple information on land cover and easily measurable landscape metrics to identify a gradient of landscape modification for the model region of Lower Saxony, Germany. We linked the gradient of landscape modification intensity to richness patterns of eight groups of vascular plants that were aggregated using residence and threat status and habitat preferences of the species.

K-means cluster analysis revealed a gradient of six landscape modification levels ranging from highly-fragmented urban landscapes to little-fragmented landscapes with large proportions of forest. Richness of all species groups differed significantly along the gradient. For the total and native species pools, for forest, low-nutrient indicating and threatened plants richness peaked at both ends of the modification intensity gradient. For neophytes, urban plants and high-nutrient indicators, richness decreased along the gradient from urbanised to less modified landscapes.

Due to its simplicity and coarse scale, our approach may easily be applied to other study areas without comprehensive data on land cover. Furthermore, following proper interpretation by experts, it can be used to identify and preselect priority landscapes for nature conservation planning.

Keywords

Landscape metrics; diversity; hemeroby; landscape naturalness; neophytes; urbanisation

Introduction

Over recent decades, tremendous man-induced changes in land cover, increasing landscape fragmentation, degradation and intensification have occurred in landscapes all over Europe (BASTIAN & BERNHARDT, 1993; MEEUS, 1995; STOATE et al., 2001; MANDER et al., 2004; FOLEY et al., 2005; WESCHE et al., 2012). Such landscape modifications have already led to large losses in (semi-)natural vegetation, biodiversity and ecosystem functions (VEAC, 2010) and are expected to cause even more losses in the future (SALA et al., 2000). Researchers have developed ideas on 'landscape modification' (FORMAN & GODRON, 1986) and the related concepts of 'landscape naturalness' (MACHADO, 2004; LIIRA & SEPP, 2009; WINTER et al., 2010), 'hemeroby' (SUKOPP, 1976; KOWARIK, 1988; GRABHERR et al., 1998; STEINHARDT et al., 1999; WRBKA et al., 2004; WALZ & STEIN, 2014), 'urbanity' (HILL et al., 2002; WRBKA et al., 2004) and 'degree of culturalness' (JANSEN et al., 2009) that are used to quantify the degree of modification a landscape has experienced. For plant species, qualifying indices are available that are based on the concepts of hemeroby (i. e., estimating the degree of human influence on the plant's environment) or urbanity (i. e., strength of association with urban environments) (JALAS, 1955; KLOTZ & KÜHN, 2002). Despite the vast number of studies exploring concepts of landscape modification or investigating human influences on biodiversity, the relationship between the two has rarely been tested directly at large scales (but see NABE-NIELSEN et al., 2010 for animal populations). Systematic studies on biodiversity patterns in modified landscapes are, however, urgently needed to assess overall effects of human impact on biodiversity and to support landscape-scale conservation planning (DAILY, 2001; FISCHER & LINDENMAYER, 2007).

There are numerous approaches to quantify the magnitude of deviation from the natural state of a landscape. While some authors have based their assessment of the landscape status solely on land cover data (STEINHARDT et al., 1999), others used multiple parameters to describe the degree of hemeroby (GRABHERR et al., 1998; WINTER et al., 2010; WALZ & STEIN, 2014) or naturalness (LIIRA & SEPP, 2009; WINTER et al., 2010) of a landscape. Most of these studies were conducted for forest habitats, but WALZ & STEIN (2014) developed an indicator of hemeroby applicable to the whole of Germany. The authors used high-resolution data on land cover and spatial information on the potential natural vegetation to derive the degree of hemeroby. They assigned the land cover categories to seven levels of hemeroby and calculated landscape hemeroby for various reference units.

Most studies on the hemeroby, naturalness or modification of a landscape or biotope are dependent on extensive field data and/or other detailed information. For many countries, such comprehensive data are not available. To avoid this problem, land cover data (either directly taken from existing digital landscape models or derived from remote sensing approaches) might prove suitable for the assessment of landscape modification and its relation to biodiversity. STEINHARDT et al. (1999, p. 2) considered data on land use/land cover as the "interface between the natural

conditions of the landscape and the influence of man". These data are "attainable within a reasonable time and with reasonable effort" (as demanded by MACHADO, 2004, p. 100). Various landscape metrics based on land cover data have been used to describe and quantify changes in landscape patterns and the composition and configuration of landscapes (LUCK & WU, 2002; YU & NG, 2006; OLSEN et al., 2007). Landscape parameters found to decrease when landscape modification increases are the amount of (semi-)natural land cover types (henceforth LCTs), as well as the average sizes of habitat patches (FORMAN & GODRON, 1986; MILLER et al., 1997; MOSER et al., 2002). As patch sizes decrease, habitats are subdivided into smaller fragments, patch numbers increase and core area sizes decrease. These processes may reduce the connectivity of patches, increase isolation between suitable habitats and enhance edge effects (HARPER et al., 2005; MASCARUA LOPEZ et al., 2006). Depending on the quality of the LCT under consideration, such landscape changes can either be interpreted as increasing the structural diversity of the landscape or increasing its subdivision (LANG & BLASCHKE, 2007).

Several studies have investigated the relationship between some of the above mentioned parameters and biota. KERR & DEGUISE (2004) and FISCHER & LINDENMAYER (2007) reported that native species benefit from the presence of natural vegetation and suggested that a decrease in the amount and size of natural habitat patches (i. e., an increase in the fragmentation of the habitat) provokes a decrease in native species richness. In contrast, many alien species are associated with human activities and commonly linked to urban habitats and to linear landscape elements. They may thus benefit from the modification of the landscape (e. g. DEUTSCHEWITZ et al., 2003; HONNAY et al., 2003; KÜHN et al., 2003). Generally, high habitat heterogeneity was found to be linked to large numbers of both native and alien species (DEUTSCHEWITZ et al., 2003).

As FORMAN & GODRON (1986, p. 286) pointed out, it is "neither possible nor useful" to separately consider the various types of human influence when evaluating the status of a landscape. Instead, the combined effects of all human influences visible at the landscape scale should be taken into account. In doing so, landscapes can be arranged along a 'landscape modification gradient'. In their well-known text on landscape ecology, FORMAN & GODRON (1986) proposed a 5-level concept on landscape modification ranging from natural to urban landscapes. Inspired by this idea, our aim was to analogously identify a gradient of landscape modification intensity based on simple information on land cover and easily measurable landscape metrics. We aimed to link the modification gradient to richness patterns of vascular plant species that were available as presence data on latitude/longitude grid cells covering the complete study area. We used LCTs that captured large proportions of the landscape and were easy to identify using remote sensing approaches. In contrast to WALZ & STEIN (2014), our analysis does not make any a priori statements about the naturalness or hemeroby of the LCTs under consideration. In the analysis, we

used the total number of vascular plant species reported for the study area as well as seven sub-groups of species aggregated using their residence and threat status as well as information on habitat preferences.

We hypothesised:

- Based on landscape metrics used to assess the composition and configuration of a landscape, clusters of landscapes with similar characteristics can be differentiated. These can be placed along a gradient of landscape modification, as proposed by FORMAN & GODRON (1986), spanning from (semi-)natural (mostly forested) to urbanised landscapes.
- The landscape clusters are meaningful for vascular plant species richness, i. e., species numbers vary significantly along the gradient of landscape modification intensity.
- Species numbers of different vascular plant species groups show different patterns along the gradient of landscape modification intensity.

Material & Methods

Study area

The study was carried out in the German federal state of Lower Saxony (c. 47,500 km², Fig. 6.1a). As landscape formation history and natural environmental configuration, colonisation history and current land use vary widely across the state, Lower Saxony harbours a large number of different habitat types.

The coastal area in the north of Lower Saxony was formed in the Holocene. It comprises the Wadden Sea, its islands and salt marshes, the estuaries of the three largest rivers of Lower Saxony (Ems, Weser and Elbe) as well as impoldered former tidal wetlands. The latter are today mainly covered by grasslands, croplands and urban areas (DRACHENFELS, 2010).

The lowlands were formed during the Pleistocene and harbour various types of landscapes, ranging from intensively managed agricultural landscapes in the west to wooded landscapes in the east.

The uplands in the south have Mesozoic and Palaeozoic bedrock and display high landscape heterogeneity. The landscape of the less elevated parts of this area is characterised by a mosaic of croplands, grasslands and various forests, while the higher elevations in the southeast (Harz Mountains) are largely covered by forests.

The largest cities of Lower Saxony are Hanover and Braunschweig, both situated at the border of the lowland and upland areas, followed by Oldenburg in the northern lowlands and Osnabrück in the western uplands.

From the coast in the north to the Harz Mountains in the southeast, elevation ranges from 0–971 m a.s.l.. From the northwest to the east, the climate gradually changes

from oceanic (mean annual precipitation up to 900 mm) to suboceanic-subcontinental (c. 550 mm; NLfB, 1997).

Lower Saxony was selected as model region because of its high landscape diversity and because the area is covered by a completed mapping program on the distribution of vascular plants with a resolution of c. 30 km² (GARVE, 2007).

Landscape composition and configuration

Following the grid system of the topographical maps, the study area was subdivided into a regular grid of 1739 cells sized c. 5.5 × 5.5 km² (30 km²; Fig. 6.1a). The 353 grid cells situated at the border of the state that only partially covered the study area and a further three grid cells completely covered by water bodies were omitted. Thus, a total of 1383 grid cells remained in the analysis; in the following each of these will be treated as a landscape unit.

Landscape composition and configuration were derived from a digital landscape model (DLM; LGN, 2008) of Lower Saxony. The DLM vector data were categorised in eight LCTs (traffic, urban and industrial areas, water bodies, forests, grasslands, croplands, and an additional class that summarizes all remaining landscape elements) and then transformed to raster format with a resolution of 25 m². Thirty different landscape metrics (Table 6) were calculated for each of the 1383 landscape units using Fragstats 3.3 (MCGARIGAL et al., 2002). Ten of these indices describe landscape composition; the remaining 20 indices depict different aspects of landscape configuration.

We calculated the proportion of each LCT per landscape unit (*Percentage of Landscape*, PLAND). The diversity at the landscape level was assessed using the *Shannon Diversity Index* (SHDI). The latter takes into account all LCTs present in the landscape as well as their extent and is especially sensitive to the presence of rare LCTs (MCGARIGAL et al., 2002).

The *Effective Mesh Size Index* (MESH) was used as an indicator to quantify the degree of fragmentation of a landscape unit. For its calculation, landscape elements that can act as barriers in the landscape (fragmentation geometries) have to be designated. In a previous analysis (SCHMIEDEL & CULMSEE, submitted), urban, industrial and traffic areas as well as water bodies were identified as suitable fragmentation geometries for the prediction of plant species richness. Thus, these LCTs were also used in the present study.

To capture average patch sizes, the *Area-weighted Mean Patch Size Index* (AREA_AM) was assessed. Together with the *Number of Patches* (NP), it supports assumptions about the fragmentation of a LCT and thus adds information to the general trend that can be derived from MESH. Patch complexity was evaluated using the *Area-weighted Mean Shape Index* (SHAPE_AM). Both, AREA_AM as well as SHAPE_AM were calculated on the landscape scale (i. e., averaged over all LCTs) as well as on the LCT level for forest, grassland and cropland areas, respectively; the number of patches was calculated for forest, grassland and cropland areas only.

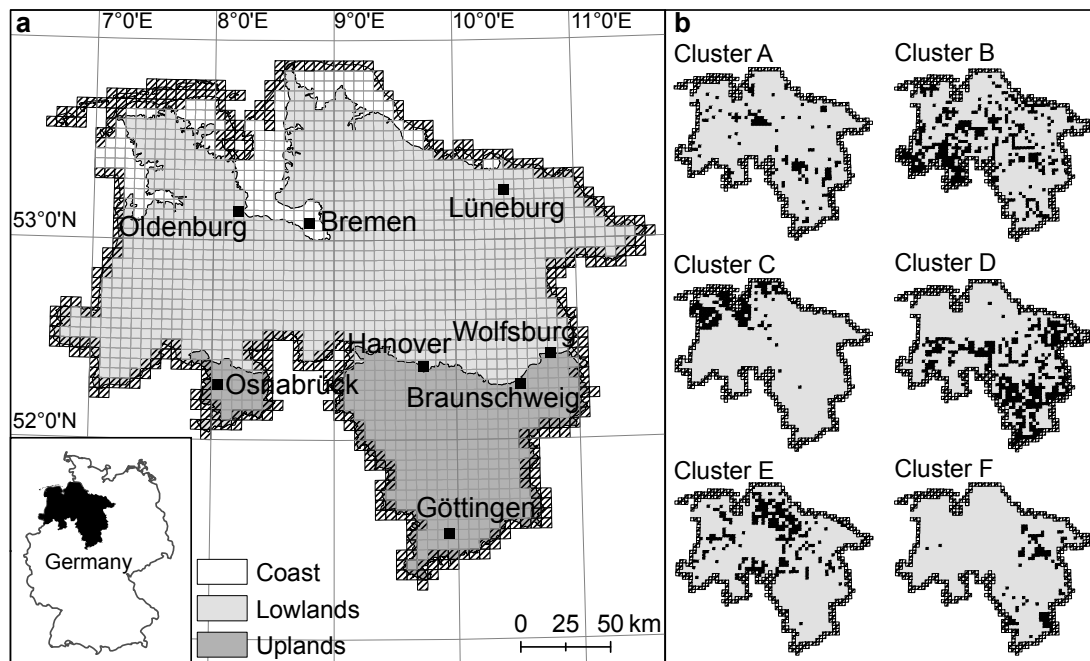


Figure 6.1: (a) The state of Lower Saxony, Germany, with its three major landscape components: coast, lowlands and uplands. The 1383 landscape units (size c. $5.5 \times 5.5 \text{ km}^2$ or 30 km^2) display the basic mapping units of the vascular plant survey program and represent the landscape units used in this analysis. Hatched grid cells were not considered in the analysis. (b) Distribution of landscape units included in clusters A to F in Lower Saxony. Maps with Transverse Mercator projection, Germany Zone 3 ("Deutsches Hauptdreiecksnetz").

Edge length was assessed using the *Total Edge Index* (TE) that sums up all edges a particular LCT shares with other LCTs.

As a measure for the isolation or rather connectivity of the patches of a LCT the *Mean Proximity Index* (PROX_MN) was assessed. It is calculated by considering size and distance of all neighbouring patches of the same type; the distance within patches that should be regarded as 'connected' has to be specified by the user ('search radius'; i. e., 100 m in the present analysis). It should be noted that only landscape connectivity and not habitat and ecological connectivity was assessed (cp. FISCHER & LINDENMAYER, 2007). TE and PROX_MN were calculated on the LCT level for forest, grassland and cropland areas, respectively.

For the LCT 'forest' the *Total Core Area Index* (TCA) and the *Mean Core Area Index* (CORE_MN) were assessed. The former measures the total amount of undisturbed core habitat after a user-specified edge buffer (i. e., 100 m in the present analysis) is eliminated (McGARIGAL et al., 2002). Analogously, CORE_MN quantifies the average amount of undisturbed core habitat in the landscape.

Table 6.1: Landscape metrics derived from land cover data (LGN, 2008) with a resolution of 25 m². Eight different land cover types (LCTs) were distinguished. Landscape metrics were calculated for 1383 landscape units covering Lower Saxony, Germany. Variables printed in bold letters were used in the cluster analysis; variables printed in normal type were highly correlated and therefore excluded from the analysis. The corresponding nomenclature and units used in Fragstats 3.3 (MCGARIGAL et al., 2002) is given. Corr., correlated variable; Transf., transformation applied to the data (log = logarithmic transformation, ~, no transformation); Mean (\pm SE), mean value across the 1383 landscape units (\pm standard error).

Variable names	Description	Fragstats nomenclature	Unit	Corr.	Transf.	Mean (\pm SE)
<i>Landscape composition</i>						
<i>prop_water</i>	Proportion of water bodies	PLAND	%	no	log	1.7 \pm 0.1
<i>prop_forest</i>	Proportion of forests	PLAND	%	no	log	21.9 \pm 0.6
<i>prop_grass</i>	Proportion of grasslands	PLAND	%	no	log	22.2 \pm 0.5
<i>prop_crop</i>	Proportion of croplands	PLAND	%	no	log	39.7 \pm 0.6
<i>prop_urban</i>	Proportion of urban areas	PLAND	%	no	log	7.2 \pm 0.2
<i>prop_industry</i>	Proportion of industrial areas	PLAND	%	no	log	2.4 \pm 0.1
<i>prop_traffic</i>	Proportion of traffic infrastructure	PLAND	%	no	log	1.7 \pm 0.0
<i>prop_others</i>	Proportion of remaining landscape elements	PLAND	%	no	log	3.3 \pm 0.1
<i>all_SHDI</i>	Shannon Diversity of all LCT (landscape diversity)	SHDI	-	no	-	1.25 \pm 0.01
<i>all_SHEI</i>	Shannon Evenness of all LCT	SHEI	-	yes	-	0.59 \pm 0.00
<i>Landscape configuration</i>						
<i>meshsize</i>	Effective Mesh Size Index (fragmentation geometries: urban, industrial and traffic area and water bodies), measure of landscape fragmentation	MESH	ha	no	-	1433.4 \pm 22.0
<i>all_AreaAm</i>	Area-weighted mean patch size averaged over all patches of all LCTs in the landscape	AREA_AM	ha	no	log	401.0 \pm 11.2

Continued on next page

Table 6.1 – Continued from previous page

Variable names	Description	Fragstats nomenclature	Unit	Corr.	Transf.	Mean (± SE)
<i>all_ShapeAm</i>	Area-weighted Mean Shape Index averaged over all patches of all LCTs in the landscape	SHAPE_AM	–	no	–	5.4±0.1
<i>for_ShapeAm</i>	Area-weighted Mean Shape Index of forest areas	SHAPE_AM	–	yes	–	2.9±0.0
<i>for_TCA</i>	Total core area of forest areas (edge depth: 100 m)	TCA	ha	yes	–	286±10.9
<i>for_AreaAm</i>	Area-weighted mean patch size of forest areas	AREA_AM	ha	yes	log	300.3±13.3
<i>for_CoreAm</i>	Area-weighted mean core area of forest areas (edge depth: 100 m)	CORE_AM	ha	yes	–	165.3±8.6
<i>for_NP</i>	Number of forest patches	NP	–	no	–	56.8±1.1
<i>for_TE</i>	Total edge of forest patches	TE	m	yes	–	89963.4±1,485.8
<i>for_ProxMn</i>	Mean Proximity Index of forest patches (search radius: 100 m)	PROX_MN	–	yes	log	3466.9±281.8
<i>grass_ShapeAm</i>	Area-weighted Mean Shape Index of grassland areas	SHAPE_AM	–	yes	–	4.3±0.1
<i>grass_AreaAm</i>	Area-weighted mean size of grassland areas	AREA_AM	ha	yes	log	130.2±6.5
<i>grass_NP</i>	Number of grassland patches	NP	–	no	–	61.6±0.7
<i>grass_TE</i>	Total edge of grassland areas	TE	m	yes	–	169293.7±3084.2
<i>grass_ProxMn</i>	Mean Proximity Index of grassland patches (search radius: 100 m)	PROX_MN	–	yes	log	4201.9±290.1
<i>crop_ShapeAm</i>	Area-weighted Mean Shape Index of cropland areas	SHAPE_AM	–	yes	–	3.8±0.1
<i>crop_AreaAm</i>	Area-weighted mean size of cropland areas	AREA_AM	ha	yes	log	289±9.5
<i>crop_NP</i>	Number of cropland patches	NP	–	no	–	43.5±0.6
<i>crop_TE</i>	Total edge of cropland areas	TE	m	yes	–	166958.7±1988.0
<i>crop_ProxMn</i>	Mean Proximity Index of cropland patches (search radius: 100 m)	PROX_MN	–	yes	log	10966.5±465.9

To improve statistical normality, the proportions of LCTs as well as the values obtained for AREA_AM and PROX_MN were log-transformed prior to the analysis. All landscape metrics were standardised to z-scores, because the range of values of the different indices strongly differed.

Landscape metrics were tested for correlation amongst each other using Spearman's rank correlation coefficient ρ . Highly correlated variables (Spearman's $\rho > |0.7|$; i. e., 15 variables, see Table 6) were excluded from the further analysis, whereas the simple landscape composition metrics were preferred over the more complex indices of landscape configuration. In total, 15 landscape metrics remained in the analysis (Table 6).

Cluster analysis

Based on the 15 standardised landscape metrics, k-means clustering was applied to pool those landscape units with a similar landscape composition and configuration. The k-means analysis groups entities according to the similarity of their attributes. It was performed using the 'kmeansruns' function from 'fpc' package (HENNING, 2014) in R version 2.15.2 (R DEVELOPMENT CORE TEAM, 2012). The function initialises the k-means algorithm several times using different starting points. The number of runs as well as the number of iterations was set to 100. The function estimates the optimal number of clusters by considering the average silhouette width criterion. The latter provides a measure of how well each landscape unit lies within the cluster it was assigned to. The index ranges between +1 (well clustered) and -1 (poorly clustered) (BROCK et al., 2008). The number of clusters to be compared was set to a minimum of 5 and a maximum of 20 clusters.

To determine the main gradient underlying the clusters, a PCA on the standardised variables was performed in R (function 'prcomp' in the 'stats' package). Clusters were sorted by the variable with most influence on the data set (i. e., prop_urban).

Species numbers per cluster

Plant species records were obtained from the data base of the vascular plant survey program of Lower Saxony (NLWKN 1982-2003) that holds information about the presence/absence of all vascular plant species occurring in the aforementioned 30 km² landscape units (GARVE, 2007).

In the analysis, we used the total number of 1594 vascular plant species reported for the study area as well as seven sub-groups of species aggregated according to their residence status, their threat status and their association with particular habitats (Table 6.2). Threatened species were identified using the Red List of Lower Saxony (GARVE, 2004). Native species were distinguished from established neophytes (KLOTZ et al., 2002). Forest species (SCHMIDT et al., 2011) were selected because they are habitat specialists that usually do not occur outside of forests. Among the eight LCTs

Table 6.2: Total number of vascular plant species of Lower Saxony, Germany, and the number of species of seven sub-groups distinguished by residence status, threat status and habitat preferences. Species groups were overlapping to some extent (cp. Table A.8 in Appendix A).

No.	Species group	Definition	Numbers of species
<i>Total vascular plant flora</i>			
1	All species	Total number of plant species recorded for Lower Saxony ^{a,b}	1594
<i>Residence status</i>			
2	Native species	Species native to Lower Saxony ^{b,c}	1297
3	Established neophytes	Species not native to Lower Saxony, established 16th century or later ^{b,c}	143
<i>Threat status</i>			
4	Threatened species	Red-listed species (threat status 1, 2, 3, G, R) ^d	538
<i>Habitat preferences</i>			
5	Forest species	Species linked to forest habitats ^e	197
6	Urban species	Species linked to urban habitats ^f	68
7	EIV species N 1-3	Species of nutrient-poor sites ^g	474
8	EIV species N 7-9	Species of nutrient-rich sites ^h	306

^a NLWKN (1982-2003); ^b GARVE (2007); ^c KLOTZ et al. (2002); ^d GARVE (2004); ^e Categories 1.1 and 1.2, SCHMIDT et al. (2011); ^f "Urbanity", categories 4 and 5, KLOTZ et al. (2002); ^g Ellenberg indicator value for nutrients (N) 1-3, BfN (2013); ^h Ellenberg indicator value for nutrients (N) 7-9, BfN (2013).

of our analysis, we considered forests representing areas least influenced by humans. Following KLOTZ et al. (2002), a group of urban species linked to settlements was defined (urbanity index 4 and 5). As eutrophication is one of the most important threats for temperate European ecosystems (DISE, 2011), two further species groups were assembled based on their association with nutrient-rich (Ellenberg Indicator Values, EIV N 7–9) and nutrient-poor habitats (EIV N 1–3), respectively (BfN, 2013). Species groups were overlapping to some extent (see Table A.8 in Appendix A). Subspecies were pooled to the corresponding species and species to the relevant aggregate of species if these were present in the data base.

Mean numbers of the total number of species reported for Lower Saxony and of the seven sub-groups were calculated per cluster. To test for significant differences between species numbers of the different clusters, a non-parametric Kruskal-Wallis test was performed in R. Multiple comparisons were made using the 'pgirmess' package (GIRAUDOUX, 2013) at a 5% significance level.

Results

Clusters of landscape modification intensity

K-means clustering of the 1383 landscape units resulted in six clusters of varying size. The smallest cluster (F) comprised 90 and the largest cluster (D) 389 landscape units (Table 6.3). The average silhouette width value of 0.16 obtained for our six clusters indicated a higher similarity within clusters than between neighbouring clusters. Although this value was relatively low, the clusters were nevertheless well interpretable.

The results of PCA (Table 6.4) suggested that the most important landscape metric was the proportion of urban areas (*prop_urban*). Further variables highly correlated with PCA axis 1 were the *Effective Mesh Size Index* (*meshsize*) and the *Area-Weighted Mean Patch Size Index* averaged over all patches of all LCTs in the landscape units (*all_AreaAm*).

The six clusters showed the following features (see Table 6.3):

Cluster A contained only 131 landscape units (9 % of the total number of investigated landscape units) and was characterised by an extremely high landscape fragmentation (very low *meshsize*, low area-weighted mean patch size). Compared to the other clusters, the patches in this cluster showed very high landscape diversity (*all_SHDI*). Most parts of these landscapes were made up by urban-industrial and traffic areas, thus the landscape units included mainly large cities (Fig. 6.1b).

Cluster B, the second largest cluster in the analysis (28 % of the landscapes units), was characterised by medium to high landscape fragmentation (low *meshsize*, low area-weighted mean patch size) and medium to high landscape diversity. All LCTs occurred at medium proportions. Forests, grasslands and croplands were each represented by a large number of patches suggesting relatively small but numerous patches (i. e., highly fragmented). Landscape units of this cluster were dispersed all over Lower Saxony (Fig. 6.1b).

Cluster C, representing 11 % of the landscape units, was characterised by numerous grassland patches, very large proportions of grassland and water bodies and low to very low proportions of forest and cropland. Landscape diversity as well as landscape fragmentation occurred at medium magnitude. The complexity of patches in the landscape was very high. Landscape units of this cluster were – with only few exceptions – mainly located in coastal and subcoastal areas and in the north-western lowlands (Fig. 6.1b).

Cluster D, the largest cluster in the analysis (28 % of the landscapes units), was dominated by croplands that on average covered more than 55 % of the landscape. Forests occurred at medium proportions, while the proportions of urban-industrial and traffic areas as well as grasslands were small. The degree of fragmentation and landscape diversity were generally medium to low. Landscape units of this cluster were mainly distributed in the uplands and the eastern and central lowlands (Fig. 6.1b).

Table 6.3: Characterisation of the landscapes in the six clusters (mean value of the variable per cluster \pm SE) found for 1383 landscape units in Lower Saxony, Germany. *n*, number of landscape units in the particular cluster. Superscript letters indicate significant differences between the clusters ($p \leq 0.05$, non-parametric multiple Kruskal-Wallis test). Non-standardized values are given; mean values of the standardized variables that were used in the analysis are given in Table A.7 in Appendix A.

Variable names	Description	A	B	C	D	E	F
<i>n</i>		131	383	148	389	242	90
Landscape composition							
prop_water	Proportion of water bodies	3.6 \pm 0.6 ^a	1.3 \pm 0.1 ^b	5.2 \pm 0.6 ^a	0.6 \pm 0.0 ^c	1.1 \pm 0.1 ^d	0.6 \pm 0.1 ^c
prop_forest	Proportion of forests	10.8 \pm 1.0 ^a	19.1 \pm 0.6 ^b	1.6 \pm 0.2 ^c	25.9 \pm 0.9 ^d	19.2 \pm 0.9 ^b	73.1 \pm 1.7 ^e
prop_grass	Proportion of grasslands	19.6 \pm 1.4 ^a	20.6 \pm 0.6 ^b	60.7 \pm 1.5 ^c	9.5 \pm 0.3 ^d	29.6 \pm 0.9 ^e	5.0 \pm 0.4 ^f
prop_crop	Proportion of croplands	26.6 \pm 1.5 ^a	44.7 \pm 0.7 ^b	21.7 \pm 1.4 ^{ac}	55.8 \pm 0.9 ^d	34 \pm 0.8 ^e	12.3 \pm 1.5 ^c
prop_urban	Proportion of urban areas	20.8 \pm 0.9 ^a	8.5 \pm 0.2 ^b	6.4 \pm 0.3 ^c	5.0 \pm 0.1 ^d	4.0 \pm 0.1 ^e	1.7 \pm 0.2 ^f
prop_industry	Proportion of industrial areas	9.0 \pm 0.6 ^a	2.2 \pm 0.1 ^b	1.1 \pm 0.1 ^c	1.1 \pm 0.1 ^c	2.9 \pm 0.4 ^d	0.7 \pm 0.1 ^e
prop_traffic	Proportion of traffic infrastructure	5.1 \pm 0.3 ^a	1.9 \pm 0.0 ^b	1.5 \pm 0.1 ^c	1.2 \pm 0.0 ^d	0.9 \pm 0.0 ^e	0.4 \pm 0.0 ^f
prop_others	Proportion of remaining landscape elements	4.4 \pm 0.3 ^a	1.8 \pm 0.1 ^b	1.9 \pm 0.2 ^b	1.0 \pm 0.0 ^c	8.4 \pm 0.5 ^d	6.1 \pm 1.3 ^b
all_SHDI	Shannon Diversity Index of all LCT	1.57 \pm 0.01 ^a	1.38 \pm 0.01 ^b	1.05 \pm 0.02 ^c	1.10 \pm 0.01 ^c	1.39 \pm 0.01 ^b	0.77 \pm 0.03 ^d
Landscape configuration							
meshsize	Effective Mesh Size Index	329.1 \pm 19.5 ^a	1042.2 \pm 27.0 ^b	1243 \pm 53.7 ^b	1761.2 \pm 36.2 ^c	1841.7 \pm 43.4 ^c	2504.4 \pm 62.3 ^d
all_AreaAm	Area-weighted Mean Patch Size Index	174.2 \pm 11.1 ^a	182.8 \pm 4.9 ^a	450.6 \pm 26.7 ^b	542 \pm 17.7 ^c	253 \pm 8.8 ^d	1367 \pm 72.9 ^e
all_ShapeAm	Area-weighted Mean Shape Index	7.5 \pm 0.3 ^a	4.9 \pm 0.1 ^b	8.5 \pm 0.2 ^a	4.9 \pm 0.1 ^{bc}	4.6 \pm 0.1 ^{bc}	4.4 \pm 0.1 ^c
for_NP	Number of forest patches	43.1 \pm 2.7 ^a	84.3 \pm 2.0 ^b	17.6 \pm 1.8 ^c	52.1 \pm 1.8 ^a	65.2 \pm 1.9 ^d	22.4 \pm 1.7 ^c
grass_NP	Number of grassland patches	67.4 \pm 2.0 ^a	81.8 \pm 1.1 ^b	46.4 \pm 1.6 ^c	56.5 \pm 1.0 ^d	55.6 \pm 1.0 ^d	30.1 \pm 1.4 ^e
crop_NP	Number of cropland patches	42.1 \pm 1.8 ^a	58.3 \pm 1.0 ^b	54.0 \pm 2.3 ^c	30.3 \pm 0.5 ^d	46.2 \pm 0.9 ^{ac}	14.5 \pm 1.1 ^e

Cluster E, including 17% of the landscape units, was characterised by a mosaic of croplands (c. 35%), grasslands (c. 30%), forests (c. 20%) and landscape elements subsumed under "others" (c. 9%). Urban-industrial and traffic areas were present at small proportions. The degree of fragmentation and landscape diversity were generally medium to low. Landscape units of this cluster were – with one exception – all located in the lowlands (Fig. 6.1b).

Cluster F, the smallest cluster in the analysis, containing 90 grid cells (7% of the landscapes units), was characterised by large proportions of forests and small proportion of all other LCTs. The values obtained for the *Effective Mesh Size Index* and area-weighted mean patch sizes were very high, thus the landscape fragmentation was low. Landscape diversity was very low. The number of grassland and cropland but also of forest patches was small. The latter in combination with the large proportion of forests suggested the occurrence of large coherent forests. These had a highly complex shape. The landscape units of this cluster were mainly situated in the Harz and Solling Mountains in the southeast of Lower Saxony as well as in the Luneburg Heath, which is part of the lowlands (Fig. 6.1b).

Species richness patterns along the landscape modification gradient

In all species groups, species richness differed significantly between the clusters (Fig. 6.2, Table A.9 in Appendix A). Richness of the total number of species reported for Lower Saxony decreased from the strongly modified landscape of cluster A to the grassland-dominated landscape of cluster C and then increased to clusters D, E and F. For the latter clusters, species numbers were highest in the cropland-dominated cluster D, followed by cluster F that was characterised by large proportions of forests (Fig. 6.2).

Corresponding patterns could be observed for the species groups of native, red-listed, forest and EIV species N1-3. While native and forest species followed the pattern of all species, the trend observed for Red List species and those species linked to nutrient-poor habitats showed a 'V shape' with the highest species numbers reported for clusters A (strongly modified landscape) and F (forest-dominated landscape) and the lowest for cluster C (grassland-dominated landscape).

The peak of species numbers in the strongly modified landscape of cluster A was repeated by the remaining species groups (neophytes, urban and EIV species N7-9), but, in contrast to the other groups, species richness more or less gradually decreased along the cluster gradient. This downward trend was only disrupted by the cropland-dominated cluster D that showed slightly higher species numbers than the surrounding clusters (Fig. 6.2).

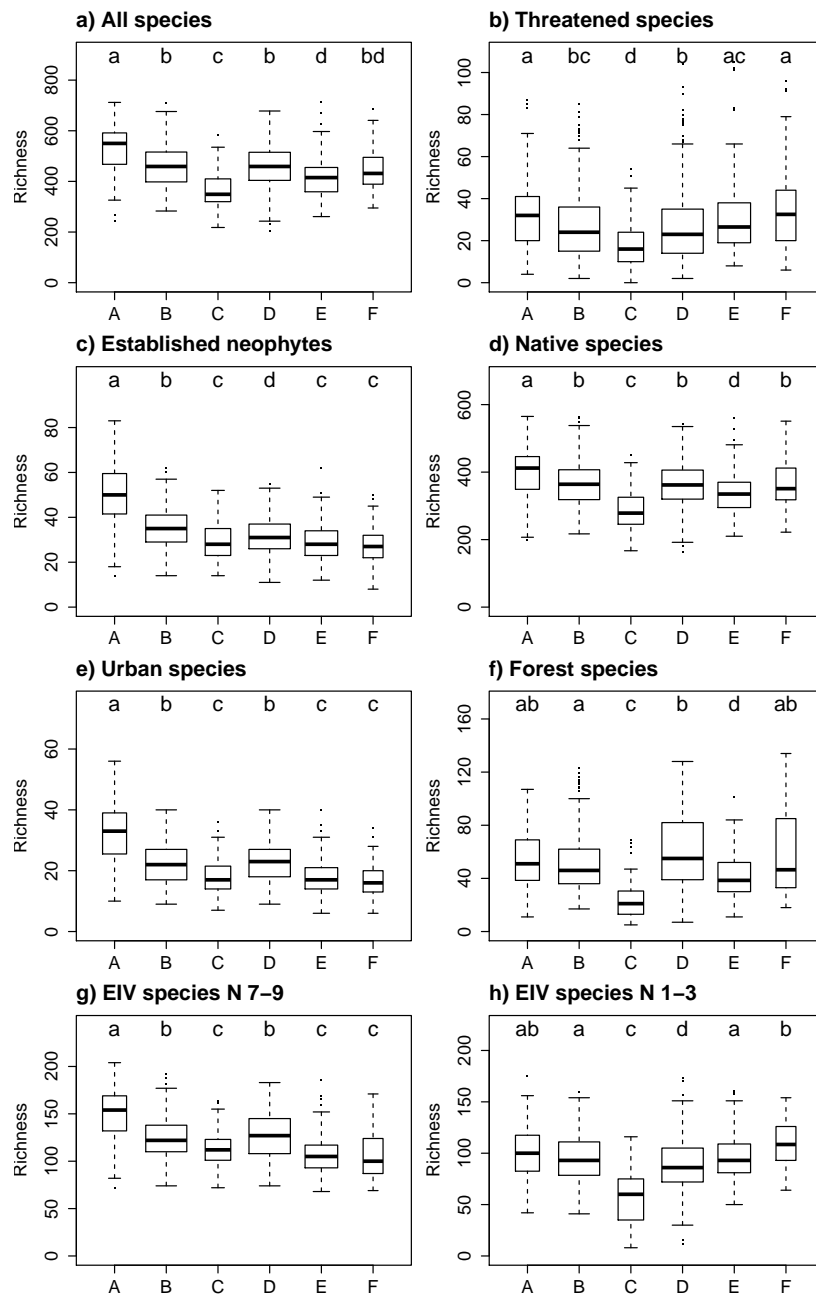


Figure 6.2: Differences in vascular plant species richness between different clusters resulting from k-means cluster analysis of 1383 landscape units (each c. 30 km²) in Lower Saxony, Germany (see also Table A.9 in Appendix A): (a) The total number of species recorded in Lower Saxony, (b) threatened species, (c) established neophytes, (d) native species, (e) urban species, (f) forest species, (g) species linked to nutrient-rich habitats (EIV species N 7-9) and (h) species linked to nutrient-poor habitats (EIV species N 1-3). Boxes are drawn with widths proportional to the square-roots of the number of observations in the groups; different letters indicate significant differences between clusters ($p \leq 0.05$, non-parametric Kruskal-Wallis test).

Table 6.4: Scores of the first ordination axis of a PCA using the standardized landscape metrics calculated for 1383 landscape units in Lower Saxony, Germany. The proportion of urban areas (prop_urban) was found to be the most important variable to describe the underlying gradient, thus clusters were sorted using this variable.

Landscape metric	Scores PCA Axis 1
prop_urban	0.37
prop_traffic	0.35
all_SHDI	0.35
prop_industry	0.29
crop_NP	0.27
grass_NP	0.26
prop_grass	0.21
prop_water	0.20
all_ShapeAm	0.11
for_NP	0.09
prop_others	0.09
prop_crop	0.04
prop_forest	-0.20
meshsize	-0.34
all_AreaAm	-0.36

Discussion

Levels of landscape modification

Our analysis revealed a gradient of landscape modification ranging from highly-fragmented and urbanised (cluster A) to little-fragmented landscapes with a large proportion of forests (cluster F). If one compares the particular clusters with the levels of landscape modification proposed by FORMAN & GODRON (1986), cluster A is an 'urban landscape' as the landscape units of this cluster are dominated by artificial, man-made elements (Table 6.5). In contrast, cluster F resembles the 'managed landscape' of FORMAN & GODRON (1986) (Table 6.5). The landscape units of this cluster were chiefly covered by forests and hosted only a very small proportion of settlements. As Lower Saxony has a very long colonization history that affected and transformed all natural habitats such as primeval forest, bogs and marshland, no 'natural landscapes' according to FORMAN & GODRON (1986) can nowadays be found.

In cluster B, all LCTs occurred at medium proportions and landscape fragmentation was quite high. The definition of 'suburban landscapes' by FORMAN & GODRON (1986) fits reasonably well to this cluster even if the landscape units were in several cases, but not necessarily, situated close to larger cities and, in the strict sense, are not 'suburbs'. Nevertheless, the authors define 'suburban landscapes' as "a town and country area with a heterogeneous, patchy mixture of residential areas, commercial centres, cropland, managed vegetation, and natural areas" (FORMAN & GODRON, 1986, p. 286), which is a good description of the situation found in the landscape units of this cluster.

The naming of the following three clusters is more difficult. Cluster C was mainly characterised by a very large proportion of grasslands and water bodies. It dominates the coastal area as well as the north-western parts of the lowlands. Its landscape is highly transformed; most natural elements, i. e., marshlands and peat bogs that had been widespread in the past have been transformed by peat digging and drainage (BEHRE, 2008). The grasslands resulting from these landscape transformations are intensively managed and interspersed with drainage ditches. Cluster D, dominated by cropland, represents those parts of the study area with intensive agricultural use. Croplands prevailed in the flatter areas while ridges were usually covered by forests. The landscape of cluster E showed a patchy mosaic of cropland, grassland and forest with relatively few settlements. All three clusters correspond to the 'cultivated landscape' of FORMAN & GODRON (1986). This shows that the simple 5-level concept of 'landscape modification' proposed by FORMAN & GODRON (1986) does not adequately differentiate our clusters when it comes to cultivated landscapes. The 10-level concept of landscape naturalness of MACHADO (2004) that allows for a more explicit differentiation seems to be more appropriate in our context. Following MACHADO (2004), the cultivated landscapes of our study may be thus better characterised as highly intervened (cluster C) to cultural assisted systems (cluster E; Table 6.5).

Species richness patterns along the landscape modification gradient

The clusters of landscape modification intensity found in this study were highly relevant for species richness patterns. Plant species numbers in all groups significantly differed between clusters. Species richness of the total number of species reported for Lower Saxony was found to peak in urban (cluster A) followed by suburban (cluster B) and managed landscapes (cluster F) whereas species richness was lowest in highly intervened landscapes (cluster C). This trend was repeated by two of the species sub-groups (native and forest species) whereas species richness of low-nutrient and threatened plants followed a 'V shaped' trend with highest species numbers in urban and managed and lowest in highly intervened landscapes. Species richness of the remaining three sub-groups (neophytes, urban and high-nutrient plants) followed a downward trend throughout from urban to managed landscapes. These results confirm our third hypothesis that species numbers of different plant species groups show different patterns along the gradient of landscape modification. The results illustrate that it is important to consider not only the entire species pool of an area but rather a number of sub-groups selected using different criteria (cp. KOWARIK, 1990).

The general peak in species numbers in urban (cluster A) and suburban landscapes (cluster B) observed for all species groups is unsurprising. Several authors have already highlighted the high species diversity of urban areas (KOWARIK, 1995; MILLER et al., 1997; HONNAY et al., 2003), which is mainly due to the higher landscape/habitat diversity (HONNAY et al., 2003; KÜHN et al., 2004) and facilitated plant immigration in urban areas (KOWARIK, 1995). In contrast, low species numbers in all groups were

found for the highly intervened landscape of cluster C. This is most likely a result of the cluster's low habitat diversity with its large proportion of transformed and intensively managed grassland areas interspersed with only few other LCTs and a low overall species diversity. The elevated species numbers in cluster D (cultural assisted to highly intervened system) reported for almost all species groups might be due to the spatial arrangement of the landscape units belonging to this cluster: Many of the landscape units are situated in the upland area and the eastern part of the lowlands with an overall larger species pool largely due to higher landscape heterogeneity (cp. SCHMIEDEL et al., 2011). Neophytes and high-nutrient plants might furthermore benefit from the large amount of intensively managed agricultural areas. The richness of forest species is also elevated in cluster D, which at first glance is surprising. This pattern may be explained by the fact that woodland patches suitable for forest plants are extant and widespread where agricultural land use was never, or is no longer, profitable.

The results show that the occurrence of non-native and urban species as well as species associated with nutrient-rich habitats is related to landscapes strongly influenced by urbanisation, fragmentation and eutrophication. For neophytes several studies confirm this trend (e. g. KOWARIK, 1990; ROY et al., 1999; KÜHN et al., 2003). By contrast, the number of native, threatened and even low-nutrient species is high in urban areas but also on the opposite side of the gradient, i. e., in the managed landscape less influenced by man. This is due to the aforementioned increased habitat diversity in urban areas that was shown to support even vulnerable species (HONNAY et al., 2003) and, regarding the other side of the landscape gradient, to the fact that largely unmodified landscapes are favourable to native plants and low-nutrient indicators (KOWARIK, 1990; FISCHER & LINDENMAYER, 2007). LENZEN et al. (2009) reported that richness of threatened species was negatively related to the amount of croplands and permanent pastures, whereas the amount of non-arable land and natural forests was positively related to threatened plant species richness. Interestingly, forest species and species linked to nutrient-poor habitats follow the same trend and show relatively high species numbers in urbanised and highly fragmented landscapes. As for forest plants, apart from remaining woodland patches, they may find refuge in urban woods and parks. Plants of nutrient-poor environments with low competitive ability are found in urban areas on gravelly and abandoned sites such as railway areas and industrial fallows. In accordance with our results, HONNAY et al. (2003) found the number of low-nutrient indicator species to be positively correlated with the amount of built-up area but did not give an explanation for this.

Table 6.5: Levels of landscape modification found for 1383 landscape units in Lower Saxony, Germany (clusters A-F) in comparison to the 'landscape modification gradient' levels by FORMAN & GODRON (1986) and the 'index of naturalness' of MACHADO (2004).

Level	Cluster ID	Description	Forman & Godron (1986)	Machado (2004)
1	Cluster A	Urban landscape, highly transformed, highly fragmented	Urban landscape	Semi-transformed to transformed system
2	Cluster B	Transformed, highly fragmented landscape with a patchy mixture of intensively managed croplands and grasslands interspersed with urban areas	Suburban landscape	Highly intervened to semi-transformed system
3	Cluster C	Transformed landscape where (semi-)natural landscape elements have been removed and water dynamics manipulated	Cultivated landscape	Highly intervened system
4	Cluster D	Transformed landscape with a large amount of intensively managed croplands (deep ploughing, fertilization); semi-natural landscape elements remained where agricultural use is not profitable	Cultivated landscape	Cultural assisted to highly intervened system
5	Cluster E	Transformed landscapes with a patchy mixture of managed systems like croplands and grasslands as well as forests	Cultivated landscape	Cultural assisted system
6	Cluster F	Managed, potentially self-maintained system (forest-dominated) with a comparatively low degree of landscape fragmentation	Managed landscape	Cultural self-maintained to cultural assisted system

Usefulness and transferability of the methodological approach

To establish the levels of landscape modification, our approach simply applies information on landscape composition and configuration in combination with k-means-clustering. Thus, it involves neither *a priori* assumptions about the hemeroby or naturalness of the LCTs under consideration (as in WALZ & STEIN, 2014) nor a subjective classification of levels of naturalness (as proposed by MACHADO, 2004). Therefore, the process of distinguishing levels of landscape modification in our concept is less susceptible to misinterpretation because only the results of the analysis and no intermediate steps need interpretation. As the levels of hemeroby of WALZ & STEIN (2014) were determined for the whole of Germany, including Lower Saxony, their results (especially those obtained for the 5 km grid) can be directly compared with those of our study. The comparison shows that the patterns derived from the two different approaches resemble each other. Both ways identify the human-dominated landscapes around the larger cities in contrast to the mainly forested areas in the southern and eastern parts of the state and designate areas under intense agricultural management.

When it comes to long-term monitoring of landscapes the concepts of WALZ & STEIN (2014) and MACHADO (2004) are superior to ours as their results may be easily compared for different time-steps as long as the same study units are used. In our approach, such results might not be directly comparable as changes in landscape composition and configuration will result in changes in the number and delimitation of clusters. Thus, our methodological approach may be applied when a 'snapshot' of the landscape status is needed and/or if due to a lack of more comprehensive information on landscape conditions more detailed assessments cannot be performed.

Similar to the concepts of WALZ & STEIN (2014) and MACHADO (2004), our approach can easily be extended to other spatial scales. The ideal scale for an analysis can be selected that fits the particular research questions and/or the spatial resolution of the available base data. The resolution of our study was restricted by the size of the landscape units adopted by the plant survey of Lower Saxony. Higher resolution through smaller landscape units and more detailed differentiation of LCTs would increase the accuracy of the assessment. With enhancing differentiation of LCTs even small-scale habitats of potential importance for nature conservation (e.g. rocky outcrops, hedgerows) could be taken into account. Nevertheless, our study showed that interpretable results can be obtained with less detailed base data.

Conclusions

Despite the fact that our approach used simple landscape metrics and was conducted at a comparatively coarse scale of c. 30 km², we were able to establish a 'gradient of landscape modification' that was shown to be meaningful for species richness. The gradient spans from highly fragmented, urbanised to less fragmented landscapes with a large proportion of forests. It was assessed without making *a priori* assump-

tions about the hemeroby or naturalness of the LCTs under consideration. The simplicity of the approach makes it transferable to regions and countries for which there is no comprehensive information on landscape conditions. The methodology can easily be extended to other spatial scales. Those parts of the landscape which are most valuable for nature conservation can be identified and preselected. It should, however, be kept in mind that the coarse scale of the analysis leaves small-scale habitats of potential importance for nature conservation undetected. We could further show that it is important to employ a set of different species groups to avoid misleading conclusions drawn from richness patterns of a single possibly inadequate group. Suitable species groups to be considered should include indicator and threatened species important for nature conservation as well as species that might pose a threat to other species or ecosystems. As high-resolution data on recent land cover (for an overview see MANAKOS & BRAUN, 2014) and the distribution of vascular plant species are becoming increasingly available in more and more countries (see Table 6 in CULMSEE et al., 2014) our approach is easily applicable to other large areas in Europe.

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7 | Synthesis

The analyses performed in the context of the present thesis aimed at improving the understanding of the relationship between species richness patterns and their underlying processes at the landscape scale with special emphasis on human influences. In particular, the analyses (a) identified and described the patterns of vascular plant species richness in the federal states of Lower Saxony and Bremen, Germany (chapter 3), (b) evaluated the adequacy of the protected area network for the protection of rare and threatened plant species in the lowland region of the study area (chapter 4) and (c) analysed the relationship between richness patterns and the human influence on the landscape (chapter 5 and 6).

The following sections recapitulate the key findings of the four main chapters of the thesis (section 7.1) and review the transferability of the methodologies applied in the thesis to other study regions (section 7.2). Furthermore, the limitations of landscape-level approaches are discussed (section 7.3). The chapter closes with concluding remarks and the open questions that build on this research (section 7.4).

7.1 Key findings and implications

7.1.1 Distinct patterns of vascular plant species richness in Lower Saxony and Bremen

The analysis presented in chapter 3 revealed a heterogeneous distribution of species richness across the study area for all six species groups considered in the study. Species numbers showed a remarkable increase from north to south and – to some extent – also from west to east. The spatial distribution of the centres of plant species richness of the different groups largely correlated with each other. However, also some interesting differences in these patterns were observed. Richness patterns of all species established in the study area largely corresponded to the patterns found for native and non-threatened species. An aggregated distribution with high overall species numbers were reported for the upland area and – with the exception of the major stream valleys of the Weser, Aller and Elbe – a much lower overall richness in the coastal and lowland area. Archaeophyte richness was particularly concentrated in the coastal area of Bremen, the Weser and Aller plains, and the loess landscapes of the lowland area¹ whereas the richness of established neophytes was highest in urban centres.

Richness patterns of threatened plants – in contrast to the patterns of all other groups – were found to be spatially highly differentiated: at the coast only the islands in the North Sea represented small hotspots, while in the lowlands relatively large hotspots were situated in the Wendland, the Luneburg Heath and the Elbe-Weser triangle north of Bremen. In the uplands, the area around Göttingen, the uplands of

¹German: *Börden*

the Weser and Leine valleys and the margins of the Harz Mountains represented well defined hotspots of Red List species.

A general trend that was observed across all species groups was that grid cells inside which urban centres were situated were found to be particularly rich in species. This relationship was especially evident for neophytes but weak for threatened species.

The presented approach can support systematic conservation planning by identifying landscape units rich in species that should receive particular attention. However, in doing so not only the total number of species but also different groups of species should be considered (cp. subsection 7.1.4). HAEUPLER & VOGEL (1999) and HAEUPLER (2000) suggest that hotspots of threatened plant species are able to reveal the "real" hotspots of vascular plant species richness and thus provide a suitable species group to be used in the determination of landscape units of high conservation value (DUELLE & OBRIST, 2003). Indeed, richness patterns of threatened species were found to be spatially much more differentiated than those of the other species groups considered in the analysis.

7.1.2 Effective 'territorial' protection of hotspots of threatened plant species

In contrast to the results of similar studies conducted for different regions of the world that identified numerous gaps in existing protected area networks (see e.g. SCOTT et al., 2001; ANDELMAN & WILLIG, 2003; DEGUISE & KERR, 2006; RODRIGUES et al., 2004), the analysis on the effectiveness of the protected area network for the conservation of threatened plant species in the lowland region of the study area confirmed a good overall coverage of the hotspots of threatened species. In particular, richness centres of the total number of threatened species were covered by nature protection areas in 85 % of the 41 geographical sub-regions considered in the analysis. The proportion of well covered hotspots increased to an effective protection in 95 % of the geographical regions when including Natura 2000 sites into the analysis. A repetition of the analyses for threatened species linked to forest habitats showed that richness centres of these species were effectively protected by nature protection areas in only 60 % of the regions; an inclusion of Natura 2000 sites increased the proportion to 81 %. However, the proven 'territorial' protection of these hotspots will only be able to preserve viable populations of the species if the protected areas are managed in an appropriate way. For many of these areas, legal regulations controlling the management of the areas are expressed very vaguely² or their development is still under way³. To effectively protect viable populations of species in the future, these regulations should be adjusted to be able to support the particular species of conservation concern. Repetitive studies in future years will then need to show if the effective 'territorial' protection is resulting in an effective species protection.

²This is especially the case for nature reserves (*Naturschutzgebiete*).

³This is mainly the case for Natura 2000 sites as these areas have been designated quite recently.

As many protected areas are isolated and small in size their effectiveness in supporting species might be limited. Therefore, areas outside reserves should also be managed in a way that supports threatened species, i. e. a "dual conservation strategy" according to JACKSON et al. (2009) should be pursued.

The methodology applied in the study was shown to be suitable to evaluate the effective 'territorial' protection of hotspots of threatened species. By transferring it to other regions with little information on the adequacy of the protected area network, it can contribute to systematic conservation planning.

7.1.3 Human impact matters: Plant species richness along gradients of landscape modification and fragmentation

The relationship between species richness patterns and human activities at the landscape scale was studied by evaluating the influence of the degree of landscape modification in general and landscape fragmentation in particular. The following two subsections highlight the key findings of chapters 5 and 6 dealing with these topics.

The *Effective Mesh Size Index* as an ecologically meaningful measure of landscape fragmentation The analysis presented in chapter 5 of the thesis demonstrated that the *Effective Mesh Size Index* (m_{eff}) can act as an ecologically meaningful measure of landscape fragmentation as long as relevant fragmentation geometries are chosen. Landscape fragmentation expressed by m_{eff} in the study area was found to vary largely between landscape units and was identified to significantly affect the richness of almost all plant species groups considered in the analysis. The size of the effect differed considerably between the groups and was largest for neophytes followed by archaeophytes. In contrast, richness patterns of native species were only slightly affected whilst those of threatened species were not affected. Instead, the variation partitioning approach for the latter species groups clearly highlighted the importance of the effect of environmental factors.

All species groups, except threatened species, showed a negative correlation with m_{eff} , i. e. richness was highest in highly fragmented and lowest in least fragmented grid cells.

In the analysis, five different m_{eff} types calculated using different fragmentation geometries were tested. Out of these, only two were found to be significant in explaining species richness patterns, namely those that used urban-industrial areas, traffic infrastructure and water bodies as fragmenting elements.

The findings of the study support the untested assumption of several researchers (e. g. JAEGER, 2000; JAEGER et al., 2008 and GIRVETZ et al., 2008) that m_{eff} is an ecologically relevant measure of landscape fragmentation. The presented approach might help future studies to determine correct fragmentation geometries to use with m_{eff} and might facilitate the unravelling of fragmentation impacts at the landscape scale. As m_{eff} in Germany is regularly assessed as one of two national indicators of land-

scape fragmentation in context of the National Strategy on Biodiversity the results obtained in the analysis are especially relevant as a contribution to future assessments.

Variation in species richness along a gradient of landscape modification Chapter 6 of this thesis applied simple information on land cover and easily measurable landscape metrics to determine a 6-level gradient of landscape modification spanning from highly fragmented urban landscapes to little fragmented ones with a large proportion of forests. Species richness of the eight groups of vascular plants considered in the analysis significantly differed along the gradient. For the entire and native species pool as well as for forest, low-nutrient indicating and threatened plants, species numbers peaked at the opposing ends of the modification gradient. For neophytes, urban plants and high-nutrient indicators, richness decreased along the gradient from high species numbers in urbanised to low numbers in less modified landscapes.

The study showed that concepts of landscape modification can be successfully linked to measures of species richness/diversity that were tested empirically here for the first time. This finding is of particular importance as concepts describing the human influence on the landscape in Germany are planned to be integrated into the nationwide monitoring system (WALZ & STEIN, 2014). In this context, knowledge on the actual relevance of such concepts for biodiversity might therefore further increase their usefulness.

The strength of the analysis is, that the assessment of the landscape modification gradient neither involved *a priori* assumptions about the hemeroby or naturalness of the land cover types under consideration (as in WALZ & STEIN, 2014) nor a subjective classification of levels of naturalness (as proposed by MACHADO, 2004). Therefore, the process of distinguishing levels of landscape modification is less susceptible to misinterpretation than the concepts of WALZ & STEIN (2014) and MACHADO (2004) because only the results of the analysis and no intermediate steps need interpretation. Due to its coarse scale, the analytical approach is transferable to other regions and countries for which there is no comprehensive information on landscape conditions. Following interpretation by experts, the methodology developed in the study allows for the identification and preselection of those parts of the landscape that are most valuable for nature conservation.

7.1.4 Overall key findings derived from the studies

In addition to the individual key findings of the four studies presented above, two further overall findings of all the analyses conducted in the present thesis can be derived that will be discussed below.

Habitat heterogeneity as a key driver of species richness in urban areas? A general emerging trend that was observed in the conducted studies on landscape fragmen-

tation and modification but also in the first study on species richness patterns, was the relationship between species richness and urban land cover: The latter together with the degree of landscape fragmentation which is usually tightly linked with the degree of urbanisation were both found to be positively associated with the richness of all species groups. The only exception being the richness of threatened species that was found to be independent of landscape fragmentation. Therefore no direct negative effects of human land use on species richness patterns could be identified. However, these results should not be used to conclude that intensive human land use increases the richness of species⁴. Instead, it seems that the elevated habitat heterogeneity of urban areas as proposed by several researchers (HAEUPLER, 1974; KÜHN et al., 2003; WANIA et al., 2006; LIPPE & KOWARIK, 2008) balances or even outbalances the assumed negative effect of landscape modification and fragmentation as long as suitable habitats are not completely removed (cp. HONNAY et al., 2003). Therefore it can be concluded that it is important to conserve the high habitat diversity of urban areas and to prevent their homogenisation.

Quality vs. quantity: avoiding drawing the wrong conclusions The analyses conducted in the framework of the thesis produced results that differed considerably between the various species groups examined in the individual studies. The richness of the groups showed different patterns according to the distribution in the study area, the coverage by protected areas and the relationship with landscape modification and fragmentation. These findings clearly emphasise the importance of considering not one single but rather an array of different species groups using various criteria. By doing so, not only quantitative but also qualitative aspects can be considered (cp. subsection 7.1.2) which are not biased to protecting mainly easily recorded or common species at the cost of a wider range of biodiversity (cp. KOWARIK, 1990; DUELLI & OBRIST, 2003; SCHETTER et al., 2013). In this context, for example, the residence or threat status of species⁵, their habitat preferences⁶ or a combination of these criteria⁷ can be used to assemble groups of species.

7.2 Transferability of the methodological approach

The analyses applied in the framework of the present thesis were conducted on a regional scale covering the area of Lower Saxony and Bremen. For this area detailed information on species occurrences and data on environmental conditions and landscape composition as well as configuration were available.

⁴Extensive management as applied in traditional cultural (pre-industrial) landscapes, in turn, is known to be beneficial for biodiversity (EEA, 2004; ANGELSTAM, 2006)

⁵As applied in SCHMIEDEL et al. (2011, 2013, submitted) and SCHMIEDEL & CULMSEE (submitted).

⁶As applied in SCHMIEDEL et al. (2013) and SCHMIEDEL et al. (submitted).

⁷As applied in SCHMIEDEL et al. (2013).

If suitable base data are at hand, the methodological approaches applied in the present thesis can be transferred to other – even larger – areas. The most important (and often the rarest) data in this context are area-wide species records. The records need to be reported on a grid- or point-basis⁸ and have to be of high quality (cp. subsection 7.3). Additionally, data on land cover and the abiotic environment are needed.

Transferability to Germany and its federal states For most of the federal states of Germany, data comparable to those used in this thesis are available that would allow a transfer. High-resolution data on land cover and topography in this thesis were taken from ATKIS-DLM and ATKIS-DEM (see section 2.2) which are both available for the whole country. Climatic and soil data can be obtained in high resolution for every federal state from the German Weather Service and the regional authorities, respectively. Concerning species records, most federal states maintain their own mapping projects (see Table 7.1). The resolutions of these mapping approaches are similar to that used for Lower Saxony or even higher (e.g. c. 8 km² grid cells for North Rhine-Westphalia, Saxony and Thuringia), the latter enabling even more detailed assessments. However, the completeness and reliability of the floristic datasets resulting from these projects would have to be tested before using them in further analyses.

With the distribution atlas of the vascular plants of Germany having been published recently (NETPHYD & BFN, 2013) reliable data on species occurrences are also available for the whole of Germany. Species data of this database are reported at the same resolution as the data for Lower Saxony (*Messtischblatt-Quadranten*, 1/4 of a topographical map). Therefore, the methodologies applied in this thesis are directly transferable to the whole country area.

Transferability to European countries Similarly to Germany, most countries in Europe maintain species mapping projects. Out of the projects listed in Table 7.1 for data acquisition only Poland uses a grid less precise than that utilised in this thesis. All other countries apply more or less equal or even higher resolutions than used in this study. As suggested for species data of the federal states of Germany, prior to an analysis, it needs to be checked whether the available information is reliable and complete.

Additional information on land cover⁹ and environmental conditions should be available for most European countries and therefore methods might be transferred to these areas as well.

The database of the *Atlas Flora Europaea* (JALAS & SUOMINEN, 1972-2013) that covers about 20 % of the European flora (FINNISH MUSEUM OF NATURAL HISTORY, 2014) pro-

⁸Point data can be transferred to grid-based data of varying resolutions.

⁹For an overview on existing data on land cover in Europe see MANAKOS & BRAUN (2014).

Table 7.1: Overview of the largest data bases on the distribution of vascular plant species in Europe, their reference areas (coverage), resolution and number of entries. Modified after CULMSEE et al. (2014, Table 6). n/a = not available.

Coverage	Resolution	Entries	Source
Europe-wide mapping project			
Europe ^d	Grid, c. 250 km ²	n/a ^b	Atlas Flora Europaea ^c
Country-wide mapping projects in Europe			
Austria	Grid, 35 km ²	c. 1,950,000	Project Mapping the Flora of Austria ^d
Belgium	Grid, 1 km ²	> 3,300,000	Florabank1 database (LANDUYT et al., 2012)
British Isles	Grid, 10 km ²	c. 9,800,000	Botanical Society of the British Isles ^e
Denmark	Grid, 25 km ²	> 1,000,000	Dansk Botanisk Forening ^f
Finland	Grid, 0.01 km ² , 1 km ² , 100 km ²	c. 5,700,000	Kasviatlas ^g
Germany	Grid, c. 30 km ²	c. 30,000,000	Verbreitungsatlas der Farn- und Blütenpflanzen Deutschlands ^h (NETPHYD & BFN, 2013)
Netherlands	Provinces, point coordinates	> 10,000,000	FLORON, Foundation for floristic research in the Netherlands ⁱ (1989 onwards)
Poland	Grid, 100 km ²	> 4,500,000	ATPOL Atlas of distribution of vascular plants in Poland (ZAJAC & ZAJAC, 2001)
Switzerland	Grid, 1 km ²	c. 3,000,000	InfoFlora National data and information centre of the flora of Switzerland ^j
State-wide mapping projects in Germany			
Baden-Württemberg	Grid, c. 30 km ²	c. 1,330,000	State Floristic Survey ^k
Bavaria	Grid, c. 33 km ²	> 1,000,000	Bavarian Floristic Survey ^l
Lower Saxony and Bremen	Grid, c. 30 km ²	2,395,045	State Inventory Program NLWKN (1982–2003), this thesis
Mecklenburg-West Pomerania	Grid, c. 30 km ²	>1,917,600	Flora of Mecklenburg-West Pomerania ^m
North Rhine-Westphalia	Grid, 8.4 km ²	c. 2,000,000	State Inventory Program (HAEUPLER et al., 2003)
Saxony	Grid, c. 8 km ²	c. 1,536,700	Project Floristic Atlas of Saxony (HARDTKE & IHL, 2000)
Thuringia	Grid, 7.8 km ²	> 1,800,000	Project Flora of Thuringia (KORSCH et al., 2002)

^a For a complete list of countries see <http://www.luomus.fi/en/atlas-florae-europaeae-afe-distribution-vascular-plants-europe>, ^b 20% of the European flora is mapped (FINNISH MUSEUM OF NATURAL HISTORY, 2014), ^c <http://www.luomus.fi/en/atlas-florae-europaeae-afe-distribution-vascular-plants-europe>, ^d <http://www.botanik.univie.ac.at/plantchorology/atlas.htm>, ^e <http://www.bsbi.org.uk/>, ^f <http://www.botaniskforening.dk/>, ^g <http://koivu.luomus.fi/kasviatlas/>, ^h <http://www.deutschlandflora.de/>, ⁱ <http://www.floron.nl/>, ^j <http://www.infoflora.ch/>, ^k <http://www.infoflora.ch/>, ^l <http://www.flora.naturkundemuseum-bw.de/>, ^m <http://www.flora-mv.de>

vides data on species distributions for the whole of Europe on a grid of c. 250 km², i.e. a resolution much coarser than that used in the present thesis. Moreover, high-resolution data on land cover for the whole area of Europe are not as easy to obtain as for the particular countries. However, information on land cover on this scale could be derived from more coarse-scaled data available for the overall area. 'Corine Land Cover 2006' (EEA, 2013) in this context provides a remote-sensing-derived raster data set on land cover with a pixel size of 100 m × 100 m that can be used for research purposes free of charge. Information on topography and climatic conditions available on a resolution of 1 km × 1 km can be derived from Worldclim data (HIJMANS et al., 2013). Therefore, potentially suitable data are available that could form the basis of analyses similar to that applied in this thesis. Because of the large differences in data resolution the results obtained from such analyses would probably, however, vary largely from those obtained at the regional scale.

Transferability to countries outside Europe The availability of high-resolution plant records for countries outside Europe is limited. However, for some areas, e.g. the state of California, USA (THE CALFLORA DATABASE, 2014) grid-based species records are acquirable. For other states of the USA (e.g. Alabama, New York, Florida, see ALABAMA HERBARIUM CONSORTIUM, 2014; NEW YORK FLORA ASSOCIATION, 2014; UNIVERSITY OF FLORIDA, INSTITUTE FOR SYSTEMATIC BOTANY, 2014) as well as for China (YANG et al., 2013) such information are not available on a grid basis but on county-level only.

Information on species records from other countries unfortunately are very hard to obtain. The transferability of the methods applied in this thesis to areas outside of Europe is therefore limited.

7.3 Limitations of landscape-level approaches

Even if frequently shown to be very powerful, landscape-level approaches are subjected to several inaccuracies arising from the scale chosen for the particular analysis, the inclusion of biased base data and inaccuracies provoked by generalisations needed to perform the analyses. GOTELLI et al. (2009) suggest that a large number of potential sources of error are linked with each data layer included in the analysis.

7.3.1 Bias due to generalisations, data resolutions and spatial scale

The representation of 'real world' conditions and patterns in 'computer space' is always subject to necessary generalisations (i.e. delimitation and discretisation processes) that help to pool the vast amount of available information. It is important that the magnitude of generalisation applied to the data should match the processes under consideration (LANG & BLASCHKE, 2007). The ATKIS-DLM data used in the present thesis represent a good example for generalisations: The DLM defines hard

boundaries along more or less homogeneous landscape elements that in most cases can not fully reflect the real conditions, as the boundaries between many land cover types (especially natural habitats) are usually more continuous. Because the latter cannot be represented cartographically and furthermore would provoke problems in data analysis, the hard boundaries are accepted as the best approximation of reality (GREEN & HARTLEY, 2000).

Generalisations affect both the spatial and the thematic resolution of the data. The latter describes the way in which the geo-spatial data are categorised¹⁰. The spatial resolution of an analysis, in turn, is described by the extent of the study area as well as the grain size¹¹. It was shown to affect all levels of data, i. e. species, land cover and environmental data.

In this context, several studies have shown that patterns of species richness/diversity change with the scale of observation or analysis (PALMER & WHITE, 1994; HE et al., 2002; KALLIMANIS et al., 2008) and therefore "areas of high species richness at one scale may appear of low species richness at the other scale" (KALLIMANIS et al., 2008, p. 148). This particular behaviour is attributable to the fact that species richness is non-additive when aggregated across different spatial scales (HE et al., 2002).

Most studies conducted at the landscape scale make use of land cover data. Usually, landscape metrics in this context are used to quantify the composition and configuration of a landscape unit. As these indices are assumed to be strongly scale-dependent, a lot of attention has focussed on the behaviour of the metrics across changing thematic and spatial resolutions. Within these studies, the thematic resolution was found to largely affect the outcome of most landscape metrics (BAILEY et al., 2007; BUYANTUYEV & WU, 2007). Related to this, the usefulness of the indices in describing landscape characteristics at varying thematic resolutions was found to largely depend on resolution: Whilst some indices are more useful at a coarser thematic resolution others are more suitable at finer scales (BAILEY et al., 2007). Similarly, only few landscape metrics have been identified to behave consistently across different spatial resolutions (TURNER et al., 1989; SAURA & MARTINEZ-MILLÁN, 2001; WU, 2004). In this context, TURNER et al. (1989) identified an increase in grain size to be responsible for the loss of rare and small-scale land cover types in the landscape. SCHINDLER et al. (2013) studied the performance of landscape metrics as indicators of species richness of various taxonomic groups across multiple scales. They found the effects of landscape structure expressed in terms of landscape metrics on species richness to strongly depend on the spatial scale of the analysis.

Finally, spatial scale also plays a role in the analysis of environmental conditions. According to SIEFERT et al. (2012) both the extent and grain size may influence the amount of variance observed in a given environmental factor. Due to the spatial au-

¹⁰cp. subsection 2.2.2 for the categorisation of DLM data applied in the present study

¹¹i. e. the size of the smallest unit of observation, (PALMER & WHITE, 1994; TURNER, 2005)

to correlation observed in most environmental variables, their variance is supposed to increase when the extent of the study area is enlarged (WIENS, 1989; SIEFERT et al., 2012). However, the size of this effect largely varies between different environmental variables. SIEFERT et al. (2012) distinguish coarse- and fine-grained variables. The former include e. g. climate data which vary across broader scales, the latter are represented by e. g. edaphic variables that may show a large variation even within a small area.

The aforementioned examples clearly illustrate the importance of scale. However, the knowledge of this scale-dependence of landscape-level approaches can only rarely be included in research as the spatial extent and grain of an analysis are usually determined by the availability of the base data.

7.3.2 Bias due to data format

"Yes, raster is faster, but raster is vaster, and vector just seems more corrector."

C. Dana Tomlin

For environmental assessments using GIS, both vector and raster data are commonly used (WADE et al., 2003). Usually, vector data are referred to as allowing high levels of cartographic accuracy¹² (WYATT & RALPHS, 2003). In contrast, raster data are supposed to be less precise than vector data as it is difficult to "represent objects with a sufficient level of cartographic precision using raster methods, since the resolution of the raster grid needs to be prohibitively high to capture an equivalent level of detail" (WYATT & RALPHS, 2003, p. 49). Nevertheless, raster data are often used in all kinds of studies because in many cases they represent the only data source available and are faster to process (LANG & BLASCHKE, 2007). For the present thesis, several data sources were available in raster format (e. g. DEM and climate data, see Table 2.1). Furthermore, some vector data (DLM) had to be converted to raster format as the software used for the subsequent analysis (Fragstats 3.3, MCGARIGAL & CUSHMAN, 2002) was limited to this type of data. In transforming the data, an additional bias is introduced because a balance between resolution and file size has to be found: If pixel size is too large, the data set will suffer from cartographic imprecision, whilst a very high resolution will generate a file too large for computation (WYATT & RALPHS, 2003). In the present thesis the resolution of the transformed data was set to a pixel size of 25 m². This value was chosen because of hardware limitations (cp. subsection 2.2.5). However, for the analyses conducted in context of this thesis this resolution should be sufficient to capture the overall characteristics of the landscape.

However, the pixel size of a raster not only affects the precision of the data set but can also directly influence the calculation of landscape metrics (MCGARIGAL & CUSHMAN, 2002; NEEL et al., 2004). Indices that include the complexity of patches are particularly affected because of the step-wise structure of the patch margins that

¹²However, this attribute still depends on the accuracy of data acquisition.

results in the overestimation of patch complexity and edge length (LEITÃO et al., 2006). In the present study this is especially true for the *Total Edge* as well as the *Shape Index* utilised in chapter 6. The latter suffers from one further limitation: Because of the raster format the simplest shape is set to be a square even though in reality circular patches have the smallest perimeter-area relationship and thus experience fewer edge effects.

7.3.3 Bias in species records

In species record data the main problem is the reliability of these data (HAEUPLER, 2000; HONNAY et al., 2003; DIEKMANN et al., 2008). Frequently, data on species occurrences are incomplete because some areas are better inventoried than others (BARTHLOTT et al., 1996). This problem apparently may provoke misleading results. Thus, it is highly important to check data on species records prior to subsequent analyses. The database utilised in the present thesis was carefully checked for its quality by the responsible authority (SCHACHERER, 2001), so that the bias related to this particular source of error should be of minor concern.

7.4 Concluding remarks and outlook

Although landscape-level approaches suffer from the "dilemma of indicating complexity with simple measures" (DUELLI & OBRIST, 2003, p. 95) and are subject to several limitations (cp. section 7.3), they are a powerful tool to disentangle the underlying causes of species richness patterns. As many, if not most, of the processes affecting biodiversity take place at the landscape level obviously they should also be measured at this scale (MCGARIGAL & CUSHMAN, 2002; FAHRIG, 2003). However, experimental approaches on such a large scale are rarely possible (WILLIG et al., 2003; GOTELLI et al., 2009), thus statistical approaches making use of existing data sources represent the only feasible method to approach this complex topic.

By applying landscape-level approaches, the present thesis showed that patterns of vascular plant species richness in Lower Saxony and Bremen are largely influenced by anthropogenically-induced land use patterns. However, also the importance of the abiotic environment in shaping species richness patterns was revealed. All analyses were conducted using the data base of the vascular plant survey of Lower Saxony that – due to its high quality and comprehensiveness – in this context provided a solid basis for these assessments.

As already highlighted in the key findings, the individual results obtained from the four studies each contribute to landscape and systematic conservation planning by facilitating the identification and preselection of landscape units of high conservation value and concern. However, by merging the knowledge derived from the studies, an even stronger comprehensive picture of the conservation value of the landscape units can be obtained. In doing so, landscape units of high conservation value, e.g. those

hosting a large number of species but have experienced a low degree of landscape modification and fragmentation, can be identified. In contrast, landscape units of high conservation concern that are species-rich despite being under high pressure from human land use, can be located. It is then possible to assess the protection status of these located landscape units by using the methodology applied in chapter 4 and determine if further conservation measures ought to be taken.

Arising from the discussed limitations and uncertainties connected to landscape-level approaches (cp. subsection 7.3), many open research questions remain that should be examined in more detail in further studies. As the transferability of the methodologies applied in this thesis is high (cp. section 7.2), it is possible to conduct analogue studies that will further increase the understanding of the factors influencing species richness patterns at the landscape scale. The most important and interesting research questions that should be addressed by future research approaches are:

1. How do the results change if the analyses are conducted at different spatial scales (changing grain size, changing spatial extent)?
2. Are the results also observed in other regions and particularly in more natural landscapes with shorter history of cultivation?
3. How do the results change if not species richness but species composition is used as the response variable in the analyses?

Progress in terms of the first research question could be made by using point-based species records, as these can be summarised into grid data of varying resolutions. For the study region of the present thesis such data could become available in the near future as the third survey period of the vascular plant survey of Lower Saxony (2004–2014) is currently assessing point-based species occurrence data.

The second research question could be addressed by transferring the methodologies of this thesis to other federal states of Germany or even other Central European countries. To test if the results are reproducible even in landscapes with a less extensive history of cultivation, a transfer to areas outside of Central Europe, e. g. to one of the northern European countries is needed. The 'Kasviatlas' of Finland (see Table 7.1) in this context could form a suitable basis for such a continuative analysis.

Finally, the third question would be particularly interesting to examine as several researchers (HILLEBRAND et al., 2008; CROWDER et al., 2010; TSCHARNTKE et al., 2012) have suggested that changes in species composition and evenness might have even larger effects on an ecosystem than changes in species richness. However, grid-based data at higher resolution than that utilised in the present thesis would probably need to be used for such an analysis as preliminary analyses conducted in context of this thesis did not return interpretable results.

By approaching these research questions, further progress will be made in the disentanglement of the "high real-world complexity shaping [species richness] patterns" (TSCHARNTKE et al., 2012, p. 678).

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List of publications

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Appendix A

Table A.1: Bio-geographical regions and natural landscape units of Lower Saxony. Basis: PILGRIM & FRANKE (1993).

Bio-geographical regions	Natural landscape unit	No.
Coast		
1 Watten und Marschen	Emsmarschen	610
	Harburger Elbmarschen	670
	Wesermarschen	612
Lowlands		
2 Ostfriesisch-Oldenburgische Geest	Aschendorfer Emstal	604
	Bourtanger Moor	605
	Hunte-Leda-Moorniederung	600
	Oldenburger Geest	603
	Ostfriesische Geest	602
	Ostfriesische Zentralmoore	601
3 Stader Geest	Achim-Verdener Geest	630
	Hamme-Oste-Niederung	632
	Wümmeniederung	631
	Wesermünder Geest	633
	Zevener Geest	634
4 Ems-Hunte-Geest und Dümmer-Geestniederung	Bersenbrücker Land	585
	Cloppenburger Geest	593
	Delmenhorster Geest	595
	Diepholzer Moorniederung	584
	Lingener Land	586
	Nordhorn-Bentheimer Sandgebiet	580
	Ostmünsterland	540
	Plantlünner Sandebene	581
	Rahden-Diepenauer Geest	582
	Sögeler Geest (Hümmling)	592
	Syker Geest	594
	Westmünsterland	544
5 Lüneburger Heide und Wendland	Hohe Heide	640
	Jeetzel-Dumme-Lehmplatte und Arendseer Platte	861
	Lüchower Niederung	860
	Luheheide	644
	Ostheide	642

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Bio-geographical regions	Natural landscape unit	No.
	Südheide	641
	Uelzener und Bevenser Becken	643
	Untere Mittelelbe-Niederung	876
	Westaltmärkisches Waldhügelland	865
6	Weser-Aller-Flachland	
	Burgdorf-Peiner Geestplatten	623
	Drömling	625
	Hannoversche Moorgeest	622
	Loccumer Geest	628
	Mittelweser	583
	Mittleres Wesertal	583
	Obere Allerniederung	626
	Ostbraunschweigisches Flachland	624
	Thedinghäuser Vorgeest	621
	Untere Aller-Talsandebene	627
	Verdener Wesertal	620
Uplands		
7	Börden	
	Bückebergvorland	522
	Braunschweig-Hildesheimer Lössbörde	520
	Calenberger Lössbörde	521
	Großes Bruch	511
	Nördliches Harzvorland, Westteil	510
	Ostbraunschweigisches Hügelland	512
8.1	Osnabrücker Hügelland	
	Östliches Wiehengebirge	532
	Lübbecker Lössland	533
	Osnabrücker Hügelland	535
	Osnabrücker Osning	534
	Ravensberger Hügelland	531
	Westliches Wiehengebirge	536
8.2	Weser- und Leinebergland	
	Alfelder Bergland (mit Ith und Hils)	377
	Eichsfelder Becken (Goldene Mark)	374
	Fulda-Werra-Bergland	357
	Göttingen-Northeimer Wald	373
	Holzmindener Wesertal	367
	Innerstebergland	379
	Kalenberger Bergland	378
	Leine-Ilme-Senke	372
	Lipper Bergland	364
	Oberwälder Land	361
	Pyrmonter Bergland	365
	Rinteln-Hamelner Weserland	366
	Südharzer Zechsteingürtel	485
	Südliches Harzvorland	486
	Südwestliches Harzvorland	376

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Table A.1 – Continued from previous page

Bio-geographical regions	Natural landscape unit	No.
	Solling, Bramwald und Reinhardswald	370
	Sollingvorland	371
	Unteres Werratal	358
	Weserengtal von Bodenwerder	368
	Westhessische Senke	343
9 Harz	Hochharz	381
	Oberharz	380
	Unterharz	382

Table A.2: List of vascular plant species of the data base of the vascular plant survey of Lower Saxony. The table lists only those species for which occurrences are reported. Extinct species are not listed. Species were aggregated on the highest taxonomical level found in the data base, i. e. subspecies were assigned to the respective species and species were assigned to the respective aggregate if the latter was listed in the data base. 'Edge' denotes that the species is present in the grid cells placed at the border of the state only (cp. section 2.1; Natural. = naturalisation (nat = native taxa, arc = archaeophytes, neo = neophytes); RL NB = Red List status assigned by GARVE (2004), categories 1, 2, 3, G and R denote threatened taxa, category 'V' denotes prewarned taxa, '*' denotes non-threatened taxa (cp. Table 2.2); EIV N1-3/N7-9 = Plant species of nutrient-poor/nutrient-rich habitats (Ellenberg Indicator Values for nutrients 1-3 and 7-9; BFN, 2013); Urban = Plant species of urban habitats (urbanity indicator 4 and 5; KLOTZ et al., 2002); Forest = Herbaceous and shrubby plant species linked to forest habitats (forest affinity categories 1.1 and 1.2; SCHMIDT et al., 2011); Code = Code used in the data base of the NLWKN (1982-2003); Incl. = Codes of (sub-)species assigned to the particular taxon, see Table A.3 for full names of (sub-)species.

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Acer campestre</i>		nat	*					cab	
<i>Acer platanoides</i>		nat	*					cad	
<i>Acer pseudoplatanus</i>		nat	*		x			cae	
<i>Achillea collina</i>		nat	R	x				caf	
<i>Achillea millefolium</i> agg.		nat	*					eab	cag
<i>Achillea pannonica</i>		nat	R	x				aaa	
<i>Achillea ptarmica</i>		nat	*	x				cah	
<i>Acinos arvensis</i>		nat	V	x				aab	
<i>Aconitum lycoctonum</i>		nat	3		x			aac	
<i>Acorus calamus</i>		neo	*		x			cai	
<i>Actaea spicata</i>		nat	*		x		x	aad	
<i>Adonis aestivalis</i>		arc	2	x				aae	
<i>Adonis vernalis</i>		nat	2	x				aag	
<i>Adoxa moschatellina</i>		nat	*		x		x	caj	
<i>Aegopodium podagraria</i>		nat	*		x			cal	
<i>Aethusa cynapium</i>		arc	*					eac	can
<i>Agrimonia eupatoria</i>		nat	*					aah	
<i>Agrimonia procera</i>		nat	*					aai	
<i>Agrostis canina</i>		nat	*	x				cap	
<i>Agrostis capillaris</i>		nat	*					caq	
<i>Agrostis stolonifera</i> agg.		nat	*					ead	cas
<i>Agrostis vinealis</i>		nat	*	x				cau	
<i>Aira caryophylla</i>		nat	V	x				aal	
<i>Aira praecox</i>		nat	*	x				aam	
<i>Ajuga genevensis</i>		nat	3	x				aao	

Continued on next page

Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Ajuga reptans</i>		nat	*					cax	
<i>Alchemilla glaucescens</i>		nat	3	x				aap	
<i>Alchemilla gramineum</i>	x	nat	1					aas	
<i>Alchemilla plicata</i>	x	nat	2					223	
<i>Alchemilla vulgaris</i> agg.		nat	*					aaq	cay, caz, cba, cbb,cbc, cbd, cbe, cbf, cbg
<i>Alisma lanceolatum</i>		nat	*					aat	
<i>Alisma plantago-aquatica</i> agg.		nat	*					eae	cbh
<i>Alliaria petiolata</i>		nat	*		x		x	cbi	
<i>Allium angulosum</i>		nat	2	x				aau	
<i>Allium carinatum</i> ssp. <i>carinatum</i>		neo	3	x				aav	
<i>Allium oleraceum</i>		nat	*					cbj	
<i>Allium paradoxum</i>		neo	*		x		x	cbk	
<i>Allium schoenoprasum</i>		neo	*	x				aax	
<i>Allium scorodoprasum</i>		nat	3		x			aay	
<i>Allium senescens</i> ssp. <i>montanum</i>		nat	1	x				aaw	
<i>Allium ursinum</i>		nat	*		x		x	aaz	
<i>Allium vineale</i>		nat	*		x			cbm	
<i>Alnus glutinosa</i>		nat	*					cbn	
<i>Alnus incana</i>		neo	*					aba	
<i>Alopecurus aequalis</i>		nat	*		x			cbo	
<i>Alopecurus bulbosus</i>		nat	2					abb	
<i>Alopecurus geniculatus</i>		nat	*		x			cbp	
<i>Alopecurus myosuroides</i>		arc	*					cbq	
<i>Alopecurus pratensis</i>		nat	*		x			cbr	
<i>Althaea hirsuta</i>		neo	2	x				cbs	
<i>Althaea officinalis</i>	x	nat	1					abc	
<i>Alyssum alyssoides</i>		nat	2	x				abd	
<i>Amaranthus blitum</i>		arc	*					363	cbv, cby
<i>Amaranthus bouchonii</i>		neo	*					cbw	
<i>Amaranthus powellii</i> agg.		neo	*			x		eag	cbz
<i>Amaranthus retroflexus</i>		neo	*		x			cca	
<i>Amelanchier lamarckii</i>		neo	*	x				ccf	
<i>Ammophila arenaria</i>		nat	*					cci	
<i>Anacamptis pyramidalis</i>		nat	2	x				abf	
<i>Anagallis arvensis</i>		arc	*					cck	eah
<i>Anagallis foemina</i>		nat	2					abg	
<i>Anagallis minima</i>		nat	3	x				ahp	
<i>Anchusa arvensis</i>		arc	*					ccm	
<i>Anchusa officinalis</i>		arc	V					abh	
<i>Andromeda polifolia</i>		nat	3	x				abi	
<i>Anemone nemorosa</i>		nat	*				x	ccn	
<i>Anemone ranunculoides</i>		nat	*		x		x	abk	
<i>Anemone sylvestris</i>		nat	2	x				abl	
<i>Anemone x semenii</i>		nat	*					242	
<i>Angelica archangelica</i>		nat	*		x			ccp	
<i>Angelica sylvestris</i>		nat	*					ccq	
<i>Antennaria dioica</i>		nat	2	x				abm	
<i>Anthemis arvensis</i>		arc	V					abn	
<i>Anthemis cotula</i>		arc	V					abo	
<i>Anthemis tinctoria</i>		arc	*					abp	
<i>Anthericum liliago</i>		nat	2	x				abq	
<i>Anthericum ramosum</i>		nat	1	x				abr	
<i>Anthoxanthum aristatum</i>		neo	*	x				cct	
<i>Anthoxanthum odoratum</i>		nat	*					ccu	
<i>Anthriscus caucalis</i>		arc	*			x		abs	
<i>Anthriscus nitida</i>		nat	R		x		x	abt	
<i>Anthriscus sylvestris</i>		nat	*		x			ccw	
<i>Anthyllis vulneraria</i>		nat	*	x				391	ccx, ccy
<i>Apera spica-venti</i>		nat	*					ccz	
<i>Aphanes arvensis</i>		arc	*					cda	
<i>Aphanes inexpectata</i>		arc	*					cdb	
<i>Apium graveolens</i>		nat	3		x			abw	
<i>Apium inundatum</i>		nat	2	x				abx	
<i>Apium repens</i>		nat	1		x			aby	
<i>Aquilegia vulgaris</i>		nat	3				x	abz	
<i>Arabidopsis thaliana</i>		arc	*					cdc	
<i>Arabis glabra</i>		nat	V					acb	
<i>Arabis hirsuta</i> agg.		nat	*					eam	
<i>Arctium lappa</i>		arc	*		x	x		cdg	
<i>Arctium minus</i>		nat	*		x	x		cdh	
<i>Arctium nemorosum</i>		nat	*		x		x	cdi	
<i>Arctium tomentosum</i>		arc	*		x	x		cdj	
<i>Arctostaphylos uva-ursi</i>		nat	2	x				acc	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Arenaria leptoclados</i>		nat	*					cdk	
<i>Arenaria serpyllifolia</i> agg.		nat	*					ean	cdl
<i>Aristolochia clematitis</i>		neo	2		x	x		acd	
<i>Armeria maritima</i>		nat	*					393	ace, acf, cdm
<i>Armoracia rusticana</i>		nat	*		x	x		cdn	
<i>Arnica montana</i>		nat	2	x				acg	
<i>Arnoseric minima</i>		nat	2	x				ach	
<i>Aronia x prunifolia</i>		neo	*					302	
<i>Arrhenatherum elatius</i>		nat	*		x			cdo	
<i>Artemisia absinthium</i>		arc	*		x			cdp	
<i>Artemisia annua</i>		neo	*			x		cdq	
<i>Artemisia biennis</i>		neo	*			x		cdr	
<i>Artemisia campestris</i>		nat	V	x				aci	
<i>Artemisia maritima</i>		nat	*		x			acj	
<i>Artemisia vulgaris</i>		nat	*		x	x		cdt	
<i>Arum maculatum</i>		nat	*		x		x	acl	
<i>Asarum europaeum</i>		nat	*				x	cdu	
<i>Asparagus officinalis</i>		arc	*					cdv	
<i>Asperugo procumbens</i>		arc	1		x			acm	
<i>Asperula cynanchica</i>		nat	2	x				aco	
<i>Asplenium adiantum-nigrum</i>		nat	1	x				acq	
<i>Asplenium ceterach</i>		nat	1	x				aia	
<i>Asplenium ruta-muraria</i>		nat	*	x		x		acu	
<i>Asplenium scolopendrium</i>		nat	3				x	ayk	
<i>Asplenium septentrionale</i>		nat	2	x				acv	
<i>Asplenium trichomanes</i>		nat	*	x				acw	224, 206, eqq, eap
<i>Asplenium viride</i>		nat	2				x	acx	
<i>Asplenium x alternifolium</i>		nat	n/a	x				acr	
<i>Asplenium x murbeckii</i>		nat	1					act	
<i>Aster amellus</i>		nat	1	x				acy	
<i>Aster linosyris</i>		nat	1	x				acz	
<i>Aster novi-belgii</i> agg.		neo	*			x		ear	cdw, cdx, cdy, 232, 50
<i>Aster tripolium</i>		nat	*		x			ada	
<i>Astragalus cicer</i>		nat	2	x			x	adb	
<i>Astragalus danicus</i>		nat	1	x				adc	
<i>Astragalus glycyphyllos</i>		nat	*	x			x	cdz	
<i>Athyrium distentifolium</i>	x	nat	R		x			add	
<i>Athyrium filix-femina</i>		nat	*				x	cea	
<i>Atriplex glabriuscula</i>		nat	R		x			ade	
<i>Atriplex laciniata</i>	x	nat	R		x			adf	
<i>Atriplex littoralis</i>		nat	*		x			cec	
<i>Atriplex longipes</i>		nat	*		x			ced	
<i>Atriplex micrantha</i>		neo	*			x		cee	
<i>Atriplex oblongifolia</i>		neo	*					cef	
<i>Atriplex patula</i>		arc	*		x			ceg	
<i>Atriplex pedunculata</i>		nat	3		x			apo	
<i>Atriplex portulacoides</i>		nat	*		x			ceh	
<i>Atriplex prostrata</i> agg.		nat	*					eas	cei
<i>Atriplex rosea</i>		neo	*		x	x		cej	
<i>Atriplex sagittata</i>		neo	*		x			cek	
<i>Atriplex tatarica</i>		neo	*					13	
<i>Atropa bella-donna</i>		nat	*		x		x	cel	
<i>Avena fatua</i>		arc	*					cem	
<i>Azolla filiculoides</i>		neo	*		x			ceo	
<i>Baldellia ranunculoides</i>		nat	2	x				adi	
<i>Ballota nigra</i>		arc	*		x	x		adj	cep, ceq
<i>Barbarea intermedia</i>		neo	*		x			cer	
<i>Barbarea stricta</i>		nat	*		x			adk	
<i>Barbarea vulgaris</i>		nat	*					ces	300, 301, 364
<i>Bellis perennis</i>		arc	*			x		ceu	
<i>Berberis vulgaris</i>		nat	3	x				adl	
<i>Berteroa incana</i>		neo	*					cev	
<i>Berula erecta</i>		nat	*					cew	
<i>Beta vulgaris</i> ssp. <i>maritima</i>	x	nat	R		x			70	
<i>Betonica officinalis</i>		nat	3	x				adm	
<i>Betula nana</i>		nat	2	x				adn	
<i>Betula pendula</i>		nat	*					cex	
<i>Betula pubescens</i>		nat	*	x				eat	cey, cez
<i>Bidens cernua</i>		nat	*		x			cfa	
<i>Bidens connata</i>		neo	*		x			cfb	
<i>Bidens frondosa</i>		neo	*		x			cfc	
<i>Bidens radiata</i>		nat	*		x			cfg	
<i>Bidens tripartita</i>		nat	*		x			cfe	

Continued on next page

Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Biscutella laevigata</i> ssp. <i>guestphalica</i>		nat	R					adp	
<i>Bistorta officinalis</i>		nat	V					azg	
<i>Blechnum spicant</i>		nat	*	x			x	adq	
<i>Blysmus compressus</i>		nat	1	x				adr	
<i>Blysmus rufus</i>	x	nat	2					ads	
<i>Bolboschoenus maritimus</i> agg.		nat	*					398	adt, dpx
<i>Botrychium lunaria</i>		nat	2	x				adu	
<i>Botrychium matricariifolium</i>		nat	1	x				adv	
<i>Brachypodium pinnatum</i>		nat	*					cfh	
<i>Brachypodium sylvaticum</i>		nat	*				x	cfi	
<i>Brassica nigra</i>		neo	*		x			cff	
<i>Briza media</i>		nat	V	x				ady	
<i>Bromus arvensis</i>		arc	3					adz	
<i>Bromus benekenii</i>		nat	*					aea	
<i>Bromus commutatus</i>		arc	*	x			x	aeb	
<i>Bromus erectus</i>		nat	*	x				aec	
<i>Bromus hordeaceus</i> agg.		nat	*					eau	cfm
<i>Bromus inermis</i>		nat	*					cfo	
<i>Bromus racemosus</i>		nat	2					aed	
<i>Bromus ramosus</i>		nat	*				x	cfq	
<i>Bromus secalinus</i>		arc	*					cfr	
<i>Bromus sterilis</i>		arc	*					cfs	
<i>Bromus tectorum</i>		arc	*					cft	
<i>Bromus thominii</i>		nat	*	x				cfm	
<i>Bryonia alba</i>		arc	3					aee	
<i>Bryonia dioica</i>		nat	V					aef	
<i>Bunias orientalis</i>		neo	*					cfu	
<i>Bunium bulbocastanum</i>		arc	2					aei	
<i>Bupleurum falcatum</i>		nat	3	x				aej	
<i>Bupleurum longifolium</i>		nat	3				x	aej	
<i>Bupleurum rotundifolium</i>		arc	1					ael	
<i>Bupleurum tenuissimum</i>		nat	2					aem	
<i>Butomus umbellatus</i>		nat	3		x			aen	
<i>Cakile maritima</i>		nat	*		x			cfv	
<i>Calamagrostis arundinacea</i>		nat	*				x	aeo	
<i>Calamagrostis canescens</i>		nat	*					cfw	
<i>Calamagrostis epigejos</i>		nat	*					cfx	
<i>Calamagrostis phragmitoides</i>		nat	R	x				68	
<i>Calamagrostis stricta</i>		nat	R	x				aep	
<i>Calamagrostis varia</i>	x	nat	R	x				aeq	
<i>Calamagrostis villosa</i>		nat	*	x				cfy	
<i>Calla palustris</i>		nat	3					aer	
<i>Callitriche palustris</i> agg.		nat	*					eay	aes, cfz, cga, aeu, cgb, cgc
<i>Calluna vulgaris</i>		nat	*	x				cgd	
<i>Caltha palustris</i>		nat	3					cge	
<i>Calystegia sepium</i>		nat	*		x			cgg	
<i>Calystegia soldanella</i>	x	nat	1					aev	
<i>Camelina microcarpa</i>		arc	2					aex	
<i>Campanula bononiensis</i>	x	nat	1	x				aez	
<i>Campanula cervicaria</i>		nat	1					afa	
<i>Campanula glomerata</i>		nat	2	x				afb	
<i>Campanula latifolia</i>		nat	3		x		x	afc	
<i>Campanula patula</i>		nat	3					afd	
<i>Campanula persicifolia</i>		nat	*	x			x	afe	
<i>Campanula rapunculoides</i>		nat	*					cgh	
<i>Campanula rapuncululus</i>		nat	*					aff	
<i>Campanula rotundifolia</i>		nat	*	x				cgi	
<i>Campanula trachelium</i>		nat	*		x		x	afg	
<i>Capsella bursa-pastoris</i>		nat	*					cgj	
<i>Cardamine amara</i>		nat	*				x	cgk	
<i>Cardamine bulbifera</i>		nat	*				x	cgl	
<i>Cardamine dentata</i>		nat	*					cgp	
<i>Cardamine flexuosa</i>		nat	*				x	cgm	
<i>Cardamine hirsuta</i>		nat	*		x			cgn	
<i>Cardamine impatiens</i>		nat	*		x		x	cgo	
<i>Cardamine parviflora</i>		nat	R					afh	
<i>Cardamine pratensis</i> agg.		nat	*					eaz	cgq
<i>Cardaminopsis arenosa</i>		neo	*	x				cgr	
<i>Cardaminopsis halleri</i>		nat	*					afi	
<i>Cardaminopsis petraea</i>	x	nat	R	x				afj	
<i>Cardaria draba</i>		neo	*					cgs	
<i>Carduus acanthoides</i>		arc	*		x			cgt	
<i>Carduus crispus</i>		nat	*		x			ebb	

Continued on next page

Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Carduus nutans</i>		arc	*					afk	
<i>Carex acuta</i>		nat	*					cgw	
<i>Carex acutiformis</i>		nat	*					cgx	
<i>Carex appropinquata</i>		nat	2					afl	
<i>Carex aquatilis</i>		nat	3					afm	
<i>Carex arenaria</i> agg.		nat	*					ebc	egy
<i>Carex bohemica</i>		nat	1					afp	
<i>Carex brizoides</i>		nat	*	x			x	cgz	
<i>Carex canescens</i>		nat	*	x				cha	
<i>Carex caryophyllea</i>		nat	*	x				afq	
<i>Carex cespitosa</i>		nat	2					afr	
<i>Carex demissa</i>		nat	V	x				chc	
<i>Carex diandra</i>		nat	2	x				afv	
<i>Carex digitata</i>		nat	*				x	afw	
<i>Carex dioica</i>		nat	1	x				afx	
<i>Carex distans</i>		nat	V					afy	
<i>Carex disticha</i>		nat	*					chd	
<i>Carex echinata</i>		nat	V	x				afz	
<i>Carex elata</i>		nat	3					chg	
<i>Carex elongata</i>		nat	3				x	chh	
<i>Carex ericetorum</i>		nat	2	x			x	aga	
<i>Carex extensa</i>		nat	*					agb	
<i>Carex flacca</i>		nat	*					agc	
<i>Carex flava</i>		nat	3	x				agd	
<i>Carex hartmanii</i>		nat	2	x				age	
<i>Carex hirta</i>		nat	*					chi	
<i>Carex hostiana</i>		nat	1	x				agf	
<i>Carex humilis</i>		nat	3	x				agg	
<i>Carex lasiocarpa</i>		nat	3	x				agh	
<i>Carex lepidocarpa</i>		nat	2	x				agi	
<i>Carex ligerica</i>		nat	*	x				agj	
<i>Carex limosa</i>		nat	1	x				agk	
<i>Carex montana</i>		nat	*	x			x	agl	
<i>Carex muricata</i> agg.		nat	*					ebe	chf, chk, chs
<i>Carex nigra</i>		nat	*	x				chl	
<i>Carex ornithopoda</i>		nat	2	x				agn	
<i>Carex otrubae</i>		nat	*					chb	
<i>Carex ovalis</i>		nat	*	x				chm	
<i>Carex pallescens</i>		nat	*	x				ago	
<i>Carex panicea</i>		nat	3					agp	
<i>Carex paniculata</i>		nat	*					chn	
<i>Carex pauciflora</i>	x	nat	2	x				agq	
<i>Carex pendula</i>		nat	*				x	cho	
<i>Carex pilosa</i>		nat	R				x	agr	
<i>Carex pilulifera</i>		nat	*	x				chp	
<i>Carex praecox</i>		nat	3					ags	
<i>Carex pseudobrizoides</i>		nat	2	x				agt	
<i>Carex pseudocyperus</i>		nat	*					agu	
<i>Carex pulicaris</i>		nat	1	x				agv	
<i>Carex punctata</i>	x	nat	2	x				agw	
<i>Carex remota</i>		nat	*				x	chq	
<i>Carex riparia</i>		nat	*					chr	
<i>Carex rostrata</i>		nat	*	x				agx	
<i>Carex strigosa</i>		nat	*				x	cht	
<i>Carex sylvatica</i>		nat	*				x	chu	
<i>Carex tomentosa</i>		nat	2					aha	
<i>Carex trinervis</i>	x	nat	2	x				ahb	
<i>Carex umbrosa</i>		nat	2				x	ahc	
<i>Carex vesicaria</i>		nat	V					chv	
<i>Carex viridula</i>		nat	V	x				agm	agy
<i>Carex vulpina</i> agg.		nat	3					ebg	chw
<i>Carex x beckmanniana</i>		nat	*					97	
<i>Carex x elytroides</i>		nat	*					chx	
<i>Carex x involuta</i>		nat	*					228	
<i>Carex x turfosa</i>		nat	R					98	
<i>Carlina acaulis</i> ssp. <i>simplex</i>		nat	2	x				ahd	
<i>Carlina vulgaris</i>		nat	*	x				ahc	
<i>Carpinus betulus</i>		nat	*					chy	
<i>Carum carvi</i>		nat	3					ahf	
<i>Catabrosa aquatica</i>		nat	2		x			ahg	
<i>Caucalis platycarpus</i>		arc	2					ahh	
<i>Centaurea cyanus</i>		arc	*					ahi	
<i>Centaurea jacea</i>		nat	*					cic	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Centaurea montana</i>		nat	2					ahj	
<i>Centaurea pseudophrygia</i>		nat	V					ahk	
<i>Centaurea scabiosa</i>		nat	*					cif	
<i>Centaureum erythraea</i>		nat	*					ahm	
<i>Centaureum littorale</i>		nat	*	x				ahn	cih, cii
<i>Centaureum pulchellum</i>		nat	*					aho	
<i>Cephalanthera damasonium</i>		nat	*				x	ahq	
<i>Cephalanthera longifolia</i>		nat	2				x	ahr	
<i>Cephalanthera rubra</i>		nat	2				x	ahs	
<i>Cerastium arvense</i>		nat	*					cij	
<i>Cerastium brachypetalum</i>		arc	2	x				aht	
<i>Cerastium diffusum</i>	x	nat	3	x				ahu	
<i>Cerastium dubium</i>		nat	3					ahv	
<i>Cerastium glomeratum</i>		arc	*					cik	
<i>Cerastium glutinosum</i>		nat	*					ahw	
<i>Cerastium holosteoides</i>		nat	*					cil	
<i>Cerastium pumilum</i> agg.		nat	*	x				ebh	ahx
<i>Cerastium semidecandrum</i>		nat	*					ahy	
<i>Ceratocarpus claviculata</i>		nat	*				x	cin	
<i>Ceratophyllum demersum</i>		nat	*		x			cio	
<i>Ceratophyllum submersum</i>		nat	*		x			ahz	
<i>Chaenorhizum minus</i>		arc	*					ciq	
<i>Chaerophyllum aureum</i>		nat	*		x			aib	
<i>Chaerophyllum bulbosum</i>		nat	*		x			aic	
<i>Chaerophyllum hirsutum</i>		nat	*		x			cir	
<i>Chaerophyllum temulum</i>		nat	*		x		x	cis	
<i>Chelidonium majus</i>		nat	*		x	x		cit	
<i>Chenopodium album</i> agg.		arc	*					ebi	cje, cjd
<i>Chenopodium bonus-henricus</i>		nat	3		x	x		aid	
<i>Chenopodium ficifolium</i>		arc	*		x			cix	
<i>Chenopodium foliosum</i>	x	neo	*		x			ciy	
<i>Chenopodium glaucum</i>		arc	*		x			ciz	
<i>Chenopodium hybridum</i>		arc	3		x			aie	
<i>Chenopodium murale</i>		arc	3		x	x		aif	
<i>Chenopodium polyspermum</i>		nat	*		x			cjb	
<i>Chenopodium rubrum</i> agg.		nat	*					ebj	cjc
<i>Chenopodium urbicum</i>		arc	1		x	x		aig	
<i>Chenopodium vulvaria</i>		arc	1		x	x		aih	
<i>Chimaphila umbellata</i>	x	nat	1	x			x	aii	
<i>Chondrilla juncea</i>		nat	3					aij	
<i>Chrysanthemum segetum</i>		arc	*					aik	
<i>Chrysosplenium alternifolium</i>		nat	*				x	ail	
<i>Chrysosplenium oppositifolium</i>		nat	*				x	aim	
<i>Cicendia filiformis</i>		nat	2	x				ain	
<i>Cicerbita alpina</i>		nat	3		x			aio	
<i>Cichorium intybus</i>		arc	*					aip	
<i>Cicuta virosa</i>		nat	3					aiq	
<i>Circaea alpina</i>		nat	*				x	air	
<i>Circaea lutetiana</i>		nat	*		x		x	cjf	
<i>Circaea x intermedia</i>		nat	*				x	ais	
<i>Cirsium acaule</i>		nat	*	x				ait	
<i>Cirsium arvense</i>		nat	*		x			cjg	
<i>Cirsium dissectum</i>		nat	2	x				aiu	
<i>Cirsium eriophorum</i>		nat	3					aiv	
<i>Cirsium oleraceum</i>		nat	*					cjh	
<i>Cirsium palustre</i>		nat	*	x				cji	
<i>Cirsium vulgare</i>		nat	*		x			cjj	
<i>Cirsium x rigens</i>		nat	*					9	
<i>Cladium mariscus</i>		nat	2	x				aix	
<i>Claytonia perfoliata</i>		neo	*		x	x		cjk	
<i>Clematis vitalba</i>		nat	*		x			cjl	
<i>Climopodium vulgare</i>		nat	*	x				aiz	
<i>Cnidium dubium</i>		arc	2					aja	
<i>Cochlearia anglica</i>		nat	V		x			cjm	
<i>Cochlearia danica</i>		nat	*					cjn	
<i>Cochlearia officinalis</i>		nat	1					ajb	
<i>Colchicum autumnale</i>		nat	3					ajd	
<i>Conium maculatum</i>		arc	*		x	x		aje	
<i>Conopodium majus</i>		neo	*					ajf	
<i>Consolida regalis</i>		arc	3					ajh	
<i>Convallaria majalis</i>		nat	*				x	aji	
<i>Convolvulus arvensis</i>		nat	*					cjr	
<i>Conyza canadensis</i>		neo	*					cjs	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Corispermum leptopterum</i>		neo	*					ajk	
<i>Cornus mas</i>		nat	3				x	ajl	
<i>Cornus sanguinea</i>		nat	*					cju	
<i>Cornus suecica</i>		nat	1	x				ajm	
<i>Coronilla coronata</i>		nat	2	x			x	ajn	
<i>Coronopus squamatus</i>		arc	3					ajp	
<i>Corrigiola litoralis</i>		nat	3					ajq	
<i>Corydalis cava</i>		nat	*		x		x	ajr	
<i>Corydalis intermedia</i>		nat	*		x		x	ajs	
<i>Corydalis solida</i>		nat	*		x		x	ajt	
<i>Corylus avellana</i>		nat	*					cjw	
<i>Corynephorus canescens</i>		nat	*	x				cjx	
<i>Cotoneaster integerrimus</i>		nat	R	x				aju	
<i>Cotula coronopifolia</i>		neo	3		x			ajv	
<i>Crambe maritima</i>		nat	R		x			ajw	
<i>Crataegus laevigata</i> agg.		nat	*					ebl	ebm, cka, ckb
<i>Crataegus monogyna</i>		nat	*					ebo	cke, ckf
<i>Crataegus rhipidophylla</i>		nat	*				x	cjz	ebk
<i>Crataegus x macrocarpa</i>		nat	*					ckd	
<i>Crataegus x media</i>		nat	*					21	
<i>Crepis biennis</i>		arc	*					ajy	
<i>Crepis capillaris</i>		arc	*					ckg	
<i>Crepis foetida</i>		arc	2	x				ajz	
<i>Crepis mollis</i>		nat	2					aka	
<i>Crepis paludosa</i>		nat	*					ckh	
<i>Crepis praemorsa</i>		nat	1	x				akb	
<i>Crepis tectorum</i>		nat	V			x		akd	
<i>Crepis vesicaria</i> ssp. <i>taraxacifolia</i>		nat	*					akc	
<i>Cruciata laevipes</i>		nat	*		x			ckj	
<i>Cucubalus baccifer</i>		nat	2		x		x	akf	
<i>Cuscuta campestris</i>		neo	*					19	
<i>Cuscuta epithymum</i>		nat	2	x				akh	
<i>Cuscuta europaea</i>		nat	*		x			ckk	
<i>Cuscuta lupuliformis</i>		neo	*		x			aki	
<i>Cymbalaria muralis</i>		neo	*			x		akj	
<i>Cynoglossum germanicum</i>		nat	3		x		x	akk	
<i>Cynoglossum officinale</i>		nat	3		x			akl	
<i>Cynosurus cristatus</i>		nat	*					cko	
<i>Cyperus fuscus</i>		nat	3					akn	
<i>Cypripedium calceolus</i>		nat	2				x	ako	
<i>Cystopteris fragilis</i>		nat	*					akp	
<i>Cytisus scoparius</i>		nat	*					ckq	
<i>Dactylis glomerata</i>		nat	*					ckr	
<i>Dactylis polygama</i>		nat	*				x	cks	
<i>Dactylorhiza incarnata</i> ssp. <i>incarnata</i>		nat	2	x				akq	
<i>Dactylorhiza maculata</i> agg.		nat	3	x				akr	ebq
<i>Dactylorhiza majalis</i>		nat	2	x				akt	
<i>Dactylorhiza praetermissa</i>		nat	3	x				aku	
<i>Dactylorhiza sphagnicola</i>		nat	2	x				akv	
<i>Dactylorhiza x dinglensis</i>		nat	n/a					dpm	
<i>Danthonia decumbens</i>		nat	V	x				ckt	
<i>Daphne mezereum</i>		nat	*				x	akw	
<i>Datura stramonium</i>		neo	*		x			akx	
<i>Daucus carota</i>		nat	*					cku	
<i>Deschampsia cespitosa</i>		nat	*	x				ckv	
<i>Deschampsia flexuosa</i>		nat	*	x				ckw	
<i>Deschampsia setacea</i>		nat	1	x				aky	
<i>Deschampsia wibeliana</i>		nat	3					akz	
<i>Descurainia sophia</i>		arc	*					ckx	
<i>Dianthus armeria</i>		nat	3	x				ala	
<i>Dianthus carthusianorum</i>		nat	3	x				alb	
<i>Dianthus deltoides</i>		nat	3	x				alc	
<i>Dianthus gratianopolitanus</i>		nat	R	x				ald	
<i>Dianthus superbus</i>		nat	1	x				ale	
<i>Digitalis grandiflora</i>		nat	2				x	alg	
<i>Digitalis purpurea</i>		nat	*				x	cky	
<i>Digitaria ischaemum</i>		arc	*	x				ckz	
<i>Digitaria sanguinalis</i>		arc	*					alh	
<i>Diphasiastrum alpinum</i>		nat	2	x				ali	
<i>Diphasiastrum complanatum</i>		nat	2	x			x	alj	
<i>Diphasiastrum issleri</i>		nat	2	x				alk	
<i>Diphasiastrum tristachyum</i>		nat	2	x				all	
<i>Diphasiastrum zeilleri</i>		nat	2	x				alm	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Diplotaxis muralis</i>		neo	*					cla	
<i>Diplotaxis tenuifolia</i>		neo	*			x		clb	
<i>Dipsacus fullonum</i>		arc	*		x			aln	
<i>Dipsacus pilosus</i>		nat	*		x		x	cld	
<i>Draba muralis</i>		neo	*			x		clf	
<i>Drosera intermedia</i>		nat	3	x				alp	
<i>Drosera longifolia</i>		nat	1	x				alo	
<i>Drosera rotundifolia</i>		nat	3	x				alr	
<i>Drosera x obovata</i>		nat	1	x				alq	
<i>Dryopteris affinis</i>		nat	*				x	als	
<i>Dryopteris carthusiana</i>		nat	*	x			x	cli	
<i>Dryopteris cristata</i>		nat	3					alt	
<i>Dryopteris dilatata</i>		nat	*		x		x	clj	
<i>Dryopteris expansa</i>		nat	R	x			x	clk	
<i>Dryopteris filix-mas</i>		nat	*				x	cll	
<i>Echinochloa crus-galli</i>		arc	*		x			clm	
<i>Echinochloa muricata</i>		neo	*					47	
<i>Echinops sphaerocephalus</i>		neo	*		x			clp	
<i>Echium vulgare</i>		arc	*					alv	
<i>Elatine hexandra</i>		nat	2	x				alx	
<i>Elatine hydropiper</i>		nat	2	x				aly	
<i>Elatine triandra</i>		nat	2					alz	
<i>Eleocharis acicularis</i>		nat	3	x				ama	
<i>Eleocharis austriaca</i>		nat	R					ebt	
<i>Eleocharis mamillata</i>		nat	3	x				amb	
<i>Eleocharis multicaulis</i>		nat	2	x				amc	
<i>Eleocharis ovata</i>		nat	1					amd	
<i>Eleocharis palustris</i>		nat	*					clr	ebu, ebv
<i>Eleocharis quinqueflora</i>		nat	3	x				amf	
<i>Eleocharis uniglumis</i>		nat	*					amg	
<i>Elodea canadensis</i>		neo	*		x			cls	
<i>Elodea nuttallii</i>		neo	*		x			clt	
<i>Elymus athericus</i>		nat	*					clv	
<i>Elymus caninus</i>		nat	*		x		x	aaJ	
<i>Elymus farctus ssp. boreoatlanticus</i>		nat	*		x			clu	
<i>Elymus repens</i>		nat	*		x			clw	
<i>Elymus x obtusiusculus</i>	x	nat	*					71	
<i>Elymus x oliveri</i>		nat	*					ctk	
<i>Empetrum nigrum</i>		nat	*	x				clx	
<i>Epilobium angustifolium</i>		nat	*		x		x	cly	
<i>Epilobium ciliatum</i>		neo	*		x			clz	
<i>Epilobium collinum</i>		nat	3	x				cma	
<i>Epilobium hirsutum</i>		nat	*		x			cmb	
<i>Epilobium lanceolatum</i>		nat	R	x				amh	
<i>Epilobium montanum</i>		nat	*					cmc	
<i>Epilobium obscurum</i>		nat	*					cmd	
<i>Epilobium palustre</i>		nat	*	x				cme	
<i>Epilobium parviflorum</i>		nat	*					cmf	
<i>Epilobium roseum</i>		nat	V		x			cmg	
<i>Epilobium tetragonum</i>		nat	*					ebw	cmh
<i>Epipactis atrorubens</i>		nat	3	x				ami	
<i>Epipactis helleborine</i>		nat	*				x	370	cmj, 243
<i>Epipactis leptochila</i>		nat	3				x	amj	cmk, cml
<i>Epipactis microphylla</i>		nat	3				x	amk	
<i>Epipactis muelleri</i>		nat	3	x			x	aml	
<i>Epipactis palustris</i>		nat	2	x			x	amm	
<i>Epipactis purpurata</i>		nat	3					amn	
<i>Epipogium aphyllum</i>		nat	1				x	amo	
<i>Equisetum arvense</i>		nat	*	x				cmm	
<i>Equisetum fluviatile</i>		nat	*					cmn	
<i>Equisetum hyemale</i>		nat	3				x	amp	
<i>Equisetum palustre</i>		nat	*	x				cmo	
<i>Equisetum pratense</i>		nat	2	x			x	amq	
<i>Equisetum sylvaticum</i>		nat	*				x	cmp	
<i>Equisetum telmateia</i>		nat	*				x	amr	
<i>Equisetum variegatum</i>		nat	1	x				ams	
<i>Equisetum x litorale</i>		nat	*					cmq	
<i>Eragrostis albensis</i>		nat	*					46	
<i>Eragrostis minor</i>		neo	*			x		cms	
<i>Erica tetralix</i>		nat	V	x				amt	
<i>Erigeron acris</i>		nat	*	x				eby	
<i>Erigeron annuus</i>		neo	*		x			cmx	401
<i>Eriophorum angustifolium</i>		nat	V	x				amu	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Eriophorum gracile</i>		nat	1	x				amv	
<i>Eriophorum latifolium</i>		nat	1	x				amw	
<i>Eriophorum vaginatum</i>		nat	V	x				amx	
<i>Erodium ballii</i>	x	nat	*	x				cmv	
<i>Erodium cicutarium</i>		nat	*					cmz	
<i>Erophila verna</i>		nat	*	x				cnb	
<i>Erucastrum gallicum</i>		neo	*					cnd	
<i>Eryngium campestre</i>		nat	3	x				amy	
<i>Eryngium maritimum</i>	x	nat	3					amz	
<i>Erysimum cheiranthoides</i>		nat	*		x			cne	
<i>Erysimum hieracifolium</i>		nat	3		x			ana	
<i>Euonymus europaea</i>		nat	*					cnh	
<i>Eupatorium cannabinum</i>		nat	*		x			cni	
<i>Euphorbia amygdaloides</i>		nat	*				x	cnj	
<i>Euphorbia cyparissias</i>		nat	*	x				cnk	
<i>Euphorbia esula</i>		nat	*					cnl	
<i>Euphorbia exigua</i>		arc	V					anb	
<i>Euphorbia helioscopia</i>		arc	*		x			cnm	
<i>Euphorbia palustris</i>		nat	2					anc	
<i>Euphorbia peplus</i>		arc	*		x			cno	
<i>Euphorbia platyphyllos</i>		nat	2					and	
<i>Euphorbia x pseudovirgata</i>		neo	*					cnp	
<i>Euphrasia frigida</i>	x	nat	R	x				248	
<i>Euphrasia nemorosa</i> agg.		nat	3					anf	cnq, cnr
<i>Euphrasia officinalis</i> ssp. <i>rostkoviana</i>		nat	3					ang	
<i>Euphrasia stricta</i>		nat	*	x				cns	
<i>Fagus sylvatica</i>		nat	*					cnu	
<i>Falcaria vulgaris</i>		nat	*					anh	
<i>Fallopia convolvulus</i>		arc	*					dah	
<i>Fallopia dumetorum</i>		nat	*		x			daj	
<i>Fallopia japonica</i>		neo	*		x			dai	
<i>Fallopia sachalinensis</i>		neo	*		x			daq	
<i>Fallopia x bohemica</i>		neo	*					64	
<i>Festuca altissima</i>		nat	*				x	cnv	
<i>Festuca arundinacea</i>		nat	*					cnw	
<i>Festuca gigantea</i>		nat	*				x	cny	
<i>Festuca heterophylla</i>		nat	3				x	ani	
<i>Festuca ovina</i> agg.		nat	*	x				ecc	coh, cnx, cnz, coa, anj, dsx, cog, 240, ank
<i>Festuca pratensis</i>		nat	*					cob	
<i>Festuca rubra</i> agg.		nat	*					397	413, ecd, 225, coe, 54, cod
<i>Filago arvensis</i>		nat	*	x				anl	
<i>Filago minima</i>		nat	*	x				ann	
<i>Filago vulgaris</i>		nat	2	x				ano	
<i>Filipendula ulmaria</i>		nat	*					ece	coi, coj
<i>Filipendula vulgaris</i>		nat	2	x				anp	
<i>Fragaria moschata</i>		nat	3				x	anq	
<i>Fragaria vesca</i>		nat	*					cok	
<i>Fragaria viridis</i>		nat	V	x				anr	
<i>Frangula alnus</i>		nat	*					col	
<i>Fraxinus excelsior</i>		nat	*		x			com	
<i>Fritillaria meleagris</i>		arc	3					ans	
<i>Fumaria officinalis</i>		arc	*		x			ecg	coq
<i>Fumaria vaillantii</i>		arc	3					ant	
<i>Gagea bohemica</i> ssp. <i>saxatilis</i>	x	nat	R	x				anx	
<i>Gagea lutea</i>		nat	*		x		x	anu	
<i>Gagea minima</i>		nat	2		x		x	anv	
<i>Gagea pratensis</i>		nat	V					anw	
<i>Gagea spathacea</i>		nat	V		x		x	cos	
<i>Gagea villosa</i>		arc	3					any	
<i>Galanthus nivalis</i>		neo	*		x		x	cot	
<i>Galeopsis angustifolia</i>		arc	3					cov	
<i>Galeopsis bifida</i>		nat	*					cow	
<i>Galeopsis ladanum</i>		nat	1	x				anz	
<i>Galeopsis segetum</i>		nat	2	x				aoa	
<i>Galeopsis speciosa</i>		nat	V		x			aob	
<i>Galeopsis tetrahit</i>		nat	*					cox	
<i>Galinsoga ciliata</i>		neo	*		x			coy	
<i>Galinsoga parviflora</i>		neo	*		x			coz	
<i>Galium album</i>		nat	*					cpa	
<i>Galium aparine</i>		nat	*		x			cpb	
<i>Galium boreale</i>		nat	3	x				aoc	
<i>Galium glaucum</i>		nat	1	x				aod	
<i>Galium odoratum</i>		nat	*				x	aoe	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Galium palustre</i>		nat	*					cpc	ctx
<i>Galium pumilum</i>		nat	3	x				cpe	
<i>Galium rotundifolium</i>		neo	R				x	aof	
<i>Galium saxatile</i>		nat	*	x				cpf	
<i>Galium spurium</i> ssp. <i>vaillantii</i>		arc	2					aog	
<i>Galium sylvaticum</i>		nat	*				x	aoh	
<i>Galium uliginosum</i>		nat	*	x				cpg	
<i>Galium verum</i>		nat	*	x				cph	aoj
<i>Galium wirtgenii</i>		nat	2					aok	
<i>Galium x pomeranicum</i>		nat	*					23	
<i>Genista anglica</i>		nat	3	x				aol	
<i>Genista germanica</i>		nat	2	x				aom	
<i>Genista pilosa</i>		nat	3	x				aon	
<i>Genista tinctoria</i>		nat	V	x				aoo	
<i>Gentiana cruciata</i>		nat	2	x				aop	
<i>Gentiana pneumonanthe</i>		nat	2	x				aoq	
<i>Gentianella ciliata</i>		nat	3	x				aou	
<i>Gentianella germanica</i>		nat	3	x				aov	
<i>Geranium columbinum</i>		arc	*		x			cpk	
<i>Geranium dissectum</i>		arc	*					cpl	
<i>Geranium lucidum</i>		nat	R		x			aos	
<i>Geranium molle</i>		arc	*					cpm	
<i>Geranium palustre</i>		nat	*		x			aoy	
<i>Geranium pratense</i>		nat	V		x			aoz	
<i>Geranium purpureum</i>		neo	*					96	
<i>Geranium pusillum</i>		arc	*		x			cpo	
<i>Geranium pyrenaicum</i>		neo	*	x		x		cpp	
<i>Geranium robertianum</i>		nat	*		x			cpq	
<i>Geranium sanguineum</i>		nat	2	x			x	apa	
<i>Geranium sylvaticum</i>		nat	*		x			apb	
<i>Geum rivale</i>		nat	3					apc	
<i>Geum urbanum</i>		nat	*		x		x	cps	
<i>Glaucium flavum</i>	x	nat	R		x			146	
<i>Glaux maritima</i>		nat	*					apd	
<i>Glechoma hederacea</i>		nat	*		x			cpt	
<i>Glyceria declinata</i>		nat	*					cpu	
<i>Glyceria fluitans</i> +		nat	*		x			cpv	
<i>Glyceria maxima</i>		nat	*		x			cpw	
<i>Glyceria notata</i>		nat	*		x			cpx	
<i>Gnaphalium sylvaticum</i>		nat	*					cpy	
<i>Gnaphalium uliginosum</i>		nat	*					cpz	
<i>Goodyera repens</i>		nat	2	x			x	apg	
<i>Gratiola officinalis</i>		nat	2					aph	
<i>Groenlandia densa</i>		nat	2					api	
<i>Gymnadenia conopsea</i>		nat	3	x				apj	
<i>Gymnocarpium dryopteris</i>		nat	*				x	apk	
<i>Gymnocarpium robertianum</i>		nat	3	x				apl	
<i>Gypsophila muralis</i>		nat	2	x				apm	
<i>Gypsophila perfoliata</i>		neo	*					43	
<i>Gypsophila repens</i>	x	nat	R	x				apn	
<i>Gypsophila scorzonerifolia</i>		neo	*					44	
<i>Hammarbya paludosa</i>		nat	1	x				app	
<i>Hedera helix</i>		nat	*					cqb	
<i>Helianthemum nummularium</i>		nat	3					apq	apr, aps
<i>Helianthus tuberosus</i>		neo	*		x			cqe	
<i>Helichrysum arenarium</i>		nat	3	x				apt	
<i>Helictotrichon pratense</i>		nat	3	x				adg	
<i>Helictotrichon pubescens</i>		nat	*					adh	
<i>Helleborus foetidus</i>		nat	*	x			x	cqf	
<i>Helleborus viridis</i>		nat	3				x	apu	
<i>Hepatica nobilis</i>		nat	*				x	apv	
<i>Heracleum mantegazzianum</i>		neo	*		x			cqg	
<i>Heracleum sphondylium</i>		nat	*		x			cqh	
<i>Herninium monorchis</i>		nat	1	x				apw	
<i>Herniaria glabra</i>		arc	*	x		x		apx	
<i>Hesperis matronalis</i>		neo	*		x		x	cqj	
<i>Hieracium aridum</i>		nat	*					dsi	
<i>Hieracium arvicola</i>		nat	G	x				cql	
<i>Hieracium aurantiacum</i>		neo	*	x				apy	
<i>Hieracium bauhini</i>		nat	*	x				apz	
<i>Hieracium bifidum</i>		nat	2	x				aqa	
<i>Hieracium bifurcum</i>		nat	G					236	
<i>Hieracium brachiatum</i>		nat	*					cqm	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Hieracium caespitosum</i>		nat	*	x				aqc	
<i>Hieracium calodon</i>		nat	G					226	
<i>Hieracium cochleatum</i>		nat	G					dwn	
<i>Hieracium cymosum</i>		nat	3	x				aqd	
<i>Hieracium densiflorum</i>		nat	G					cqo	
<i>Hieracium diaphanoides</i>		nat	*					cqp	
<i>Hieracium flagellare</i>		nat	G					cqr	
<i>Hieracium flagelliferum</i>		nat	*					dps	
<i>Hieracium floribundum</i>		nat	G					cqs	
<i>Hieracium glaucinum</i>		nat	*	x				aqe	
<i>Hieracium glaucisetigerum</i>		nat	G					278	
<i>Hieracium glomeratum</i>		nat	G					279	
<i>Hieracium guthmickianum</i>		nat	*					7	
<i>Hieracium kalksburgense</i>		nat	G	x				79	
<i>Hieracium lachenalii</i>		nat	*	x				cqt	
<i>Hieracium lactucella</i>		nat	2	x				aqf	
<i>Hieracium laevigatum</i>		nat	*	x				cqu	
<i>Hieracium laurinum</i>		nat	*					cqv	
<i>Hieracium leptophyton</i>		nat	G					cqw	
<i>Hieracium longiscapum</i>		nat	G					dwm	
<i>Hieracium macrostolonum</i>		nat	*					195	
<i>Hieracium maculatum</i>		nat	G	x				cqx	
<i>Hieracium murorum</i>		nat	*				x	aqi	
<i>Hieracium pilosella</i>		nat	*	x				cqy	
<i>Hieracium piloselloides</i>		nat	*	x				cqz	
<i>Hieracium polymastix</i>		nat	*					218	
<i>Hieracium prussicum</i>		nat	G					212	
<i>Hieracium rothianum</i>		nat	R	x				cqq	
<i>Hieracium sabaudum</i>		nat	*	x				cra	
<i>Hieracium saxifragum</i>	x	nat	R					233	
<i>Hieracium schmidtii</i>		nat	1	x				aqg	
<i>Hieracium schultesii</i>	x	nat	1	x				crb	
<i>Hieracium stoloniflorum</i>		nat	*					219	
<i>Hieracium umbellatum</i>		nat	*	x				crc	
<i>Hieracium visianii</i>		nat	*					280	
<i>Hieracium vulgatum</i>		nat	*	x				crd	
<i>Hieracium zizianum</i>		nat	G					cre	
<i>Hierochloë odorata</i>		nat	3	x				aqj	
<i>Hippocrepis comosa</i>		nat	3	x				aqk	
<i>Hippophaë rhamnoides</i>	x	nat	*	x				crf	
<i>Hippuris vulgaris</i>		nat	3					aqi	
<i>Holcus lanatus</i>		nat	*					crh	
<i>Holcus mollis</i>		nat	*	x				cri	
<i>Holostemum umbellatum</i>		nat	V	x				aqm	
<i>Honckenya peploides</i>		nat	*		x			crj	
<i>Hordelymus europaeus</i>		nat	*				x	crk	
<i>Hordeum jubatum</i>		neo	*			x		crl	
<i>Hordeum murinum</i>		arc	*			x		crm	
<i>Hordeum secalinum</i>		nat	V					aqo	
<i>Hornungia petraea</i>		nat	1	x				aqp	
<i>Hottonia palustris</i>		nat	V					aqq	
<i>Humulus lupulus</i>		nat	*		x		x	crn	
<i>Huperzia selago</i>		nat	3				x	aqr	
<i>Hydrocharis morsus-ranae</i>		nat	V					aqs	
<i>Hydrocotyle vulgaris</i>		nat	*	x				aqt	
<i>Hymenolobus procumbens</i>		neo	*					36	
<i>Hyoscyamus niger</i>		arc	3		x			aqu	
<i>Hypericum elodes</i>		nat	2	x				aqv	
<i>Hypericum hirsutum</i>		nat	*		x		x	aqw	
<i>Hypericum humifusum</i>		nat	*	x				aqx	
<i>Hypericum maculatum</i>		nat	*	x				crp	ecm
<i>Hypericum montanum</i>		nat	3	x				aqy	
<i>Hypericum perforatum</i>		nat	*					crq	
<i>Hypericum pulchrum</i>		nat	*	x				aqz	
<i>Hypericum tetrapterum</i>		nat	*					crr	
<i>Hypochaeris glabra</i>		nat	2	x				ara	
<i>Hypochaeris maculata</i>		nat	1	x				arb	
<i>Hypochaeris radicata</i>		nat	*	x				crs	
<i>Ilex aquifolium</i>		nat	*				x	crv	
<i>Illecebrum verticillatum</i>		nat	3	x				arc	
<i>Impatiens glandulifera</i>		neo	*		x			crw	
<i>Impatiens noli-tangere</i>		nat	*				x	crx	
<i>Impatiens parviflora</i>		neo	*				x	cry	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Inula britannica</i>		nat	3					ard	
<i>Inula conyzae</i>		nat	*	x				crz	
<i>Inula germanica</i>	x	nat	R	x				are	
<i>Inula helenium</i>		neo	*					arf	
<i>Inula hirta</i>		nat	R	x				arg	
<i>Inula salicina</i>		nat	3	x				arh	
<i>Iris pseudacorus</i>		nat	*		x			csa	
<i>Iris sibirica</i>		nat	2	x				ari	
<i>Iris versicolor</i>		neo	*					csb	
<i>Isoetes lacustris</i>		nat	1	x				arj	
<i>Isolepis fluitans</i>		nat	2	x				ark	
<i>Isolepis setacea</i>		nat	3	x				arl	
<i>Jasione montana</i>		nat	*	x				arm	
<i>Juncus acutiflorus</i>		nat	*	x				cse	
<i>Juncus alpinus</i>		nat	2	x				arn	
<i>Juncus anceps</i>	x	nat	*					aro	
<i>Juncus articulatus</i>		nat	*	x				csf	
<i>Juncus bufonius</i>		nat	*					csg	
<i>Juncus bulbosus</i>		nat	*	x				csk	
<i>Juncus capitatus</i>		nat	1	x				arq	
<i>Juncus compressus</i>		nat	*					csi	
<i>Juncus conglomeratus</i>		nat	*	x				csj	
<i>Juncus effusus</i>		nat	*					csk	
<i>Juncus filiformis</i>		nat	3	x				arr	
<i>Juncus gerardii</i>		nat	*					ars	
<i>Juncus inflexus</i>		nat	*					art	
<i>Juncus maritimus</i>	x	nat	*					aru	
<i>Juncus minutulus</i>		nat	D					ecp	
<i>Juncus ranarius</i>		nat	*	x				csm	
<i>Juncus squarrosus</i>		nat	V	x				csn	
<i>Juncus subnodulosus</i>		nat	2	x				arv	
<i>Juncus tenageia</i>		nat	2					arw	
<i>Juncus tenuis</i>		neo	*					cso	
<i>Juniperus communis</i>		nat	3					arx	
<i>Kalmia angustifolia</i>		neo	*	x		x		csp	
<i>Kickxia elatine</i>		arc	2	x				ary	
<i>Kickxia spuria</i>		arc	2	x				arz	
<i>Knautia arvensis</i>		nat	*					csq	
<i>Koeleria arenaria</i>	x	nat	V	x				asa	
<i>Koeleria glauca</i>		nat	2	x				asb	
<i>Koeleria macrantha</i>		nat	3	x				asc	
<i>Koeleria pyramidata</i>		nat	V	x				css	
<i>Lactuca serriola</i>		nat	*			x		csu	
<i>Lactuca tatarica</i>		neo	*					csv	
<i>Lactuca virosa</i>		nat	3		x			ase	
<i>Lamium album</i>		arc	*		x			csw	
<i>Lamium amplexicaule</i>		arc	*		x			csx	
<i>Lamium argentatum</i>		neo	*			x		ecr	
<i>Lamium galeobdolon</i> agg.		nat	*					ecq	csz, cta
<i>Lamium maculatum</i>		nat	*		x			ctb	
<i>Lamium purpureum</i>		arc	*		x			ect	asf
<i>Lapsana communis</i>		nat	*		x			ctd	
<i>Laser trilobum</i>		nat	3	x			x	ash	
<i>Laserpitium latifolium</i>		nat	2	x				asi	
<i>Lathraea squamaria</i>		nat	V				x	ask	
<i>Lathyrus hirsutus</i>		neo	*					ctg	
<i>Lathyrus latifolius</i>		neo	*	x		x		cth	
<i>Lathyrus linifolius</i>		nat	V	x				asm	
<i>Lathyrus maritimus</i>	x	nat	R	x				asn	
<i>Lathyrus niger</i>		nat	2	x			x	aso	
<i>Lathyrus missolia</i>		neo	*					cti	
<i>Lathyrus palustris</i>		nat	2	x				asp	
<i>Lathyrus pratensis</i>		nat	*					ctj	
<i>Lathyrus sylvestris</i>		nat	*	x				asq	
<i>Lathyrus tuberosus</i>		nat	V					asr	
<i>Lathyrus vernus</i>		nat	*				x	ctl	
<i>Ledum palustre</i>		nat	2	x				ass	
<i>Leersia oryzoides</i>		nat	2		x			ast	
<i>Legousia hybrida</i>		arc	2	x				asu	
<i>Legousia speculum-veneris</i>		arc	1	x				asv	
<i>Lemna gibba</i>		nat	*		x			ctm	
<i>Lemna minor</i>		nat	*					ctn	
<i>Lemna minuta</i>		neo	*					cto	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Lemna trisulca</i>		nat	*					ctp	
<i>Lemna turionifera</i>		neo	*					28	
<i>Leontodon autumnalis</i>		nat	*					ctq	
<i>Leontodon hispidus</i>		nat	*					asw	
<i>Leontodon saxatilis</i>		nat	*					asx	
<i>Leonurus cardiaca</i>		arc	*		x	x		asy	ctr, cts
<i>Leonurus marrubiastrum</i>		nat	3		x			asz	
<i>Lepidium campestre</i>		arc	*					ata	
<i>Lepidium heterophyllum</i>		neo	*			x		ctu	
<i>Lepidium latifolium</i>		neo	*					ctv	
<i>Lepidium ruderale</i>		arc	*					cty	
<i>Leucanthemum vulgare</i> agg.		nat	*	x				ecu	cub
<i>Leucojum vernum</i>		nat	*		x			atc	
<i>Leymus arenarius</i>		nat	*					cud	
<i>Ligustrum vulgare</i>		nat	*	x				cue	
<i>Lilium bulbiferum</i>		nat	2	x				atd	ate, atf
<i>Lilium martagon</i>		nat	3					atg	
<i>Limonium vulgare</i>		nat	V					ath	
<i>Limosella aquatica</i>		nat	3	x				ati	
<i>Linaria arvensis</i>		arc	1					atj	
<i>Linaria vulgaris</i>		nat	*					cug	
<i>Lindernia dubia</i>		neo	*					atk	
<i>Linnaea borealis</i>		nat	1	x			x	atl	
<i>Linum catharticum</i>		nat	*	x				atm	
<i>Linum leonii</i>		nat	2	x				atn	
<i>Linum tenuifolium</i>		nat	1	x				ato	
<i>Liparis loeselii</i>		nat	2	x				atp	
<i>Listera cordata</i>		nat	3	x			x	atq	
<i>Listera ovata</i>		nat	*		x		x	atr	
<i>Lithospermum arvense</i>		arc	3					aeg	
<i>Lithospermum officinale</i>		nat	3					ats	
<i>Lithospermum purpureocaruleum</i>		nat	3				x	aeH	
<i>Littorella uniflora</i>		nat	2	x				att	
<i>Lobelia dortmanna</i>		nat	1	x				atu	
<i>Lolium perenne</i>		nat	*		x			cuk	
<i>Lonicera periclymenum</i>		nat	*					cum	
<i>Lonicera xylosteum</i>		nat	*				x	atw	
<i>Lotus corniculatus</i>		nat	*	x				cun	
<i>Lotus pedunculatus</i>		nat	*					cuo	
<i>Lotus tenuis</i>		nat	V					atx	
<i>Lunaria rediviva</i>		nat	V		x		x	atz	
<i>Lupinus polyphyllus</i>		neo	*					cus	
<i>Luronium natans</i>		nat	2	x				aua	
<i>Luzula campestris</i>		nat	*	x				cut	
<i>Luzula congesta</i>		nat	3					cuu	
<i>Luzula luzuloides</i>		nat	*				x	aub	238
<i>Luzula multiflora</i>		nat	*	x				cuv	
<i>Luzula pilosa</i>		nat	*				x	cuw	
<i>Luzula sudetica</i>		nat	R	x				cux	
<i>Luzula sylvatica</i>		nat	*					auc	
<i>Lycium barbarum</i>		neo	*			x		cuz	
<i>Lycopodiella inundata</i>		nat	3	x				aeu	
<i>Lycopodium annotinum</i>		nat	3	x			x	auf	
<i>Lycopodium clavatum</i>		nat	3	x				aug	
<i>Lycopus europaeus</i>		nat	*		x			cvb	
<i>Lysimachia nemorum</i>		nat	*		x		x	auh	
<i>Lysimachia nummularia</i>		nat	*					cvc	
<i>Lysimachia thyrsoflora</i>		nat	V					auj	
<i>Lysimachia vulgaris</i>		nat	*					cve	
<i>Lythrum hyssopifolia</i>		nat	1					auj	
<i>Lythrum salicaria</i>		nat	*					cvf	
<i>Maianthemum bifolium</i>		nat	*	x			x	cvh	
<i>Malus sylvestris</i>		nat	3					auk	
<i>Malva alcea</i>		arc	V		x			cvj	
<i>Malva moschata</i>		arc	*			x		cvk	
<i>Malva neglecta</i>		arc	*		x	x		cvl	
<i>Malva pusilla</i>		arc	2			x		cvm	
<i>Malva sylvestris</i> ssp. <i>sylvestris</i>		arc	*			x		cvn	
<i>Marrubium vulgare</i>		arc	1		x			aul	
<i>Matricaria discoidea</i>		neo	*		x	x		cvp	
<i>Matricaria recutita</i>		arc	*					cvq	
<i>Matteuccia struthiopteris</i>		nat	3		x		x	aum	
<i>Medicago falcata</i>		nat	*	x				cvs	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Medicago lupulina</i>		nat	*					cvt	
<i>Medicago minima</i>		nat	2	x				aun	
<i>Medicago x varia</i>		neo	*					cvu	
<i>Melampyrum arvense</i>		nat	2	x				auo	
<i>Melampyrum cristatum</i>		nat	2	x			x	aup	
<i>Melampyrum nemorosum</i>		nat	V				x	auq	
<i>Melampyrum pratense</i>		nat	*	x				cvv	
<i>Melampyrum sylvaticum</i>		nat	*	x				aur	
<i>Melica nutans</i>		nat	*	x			x	aus	
<i>Melica uniflora</i>		nat	*				x	cvw	
<i>Melilotus albus</i>		arc	*					cvx	
<i>Melilotus altissimus</i>		nat	*		x			cvy	
<i>Melilotus dentatus</i>		nat	1					aut	
<i>Melilotus officinalis</i>		arc	*	x				cwa	
<i>Melittis melissophyllum</i>		nat	1	x			x	auu	
<i>Mentha aquatica</i>		nat	*					cwb	
<i>Mentha arvensis</i>		nat	*					cwc	
<i>Mentha longifolia</i>		nat	*		x			auv	
<i>Mentha pulegium</i>		nat	2		x			auw	
<i>Mentha x verticillata</i>		nat	*					cwi	
<i>Menyanthes trifoliata</i>		nat	3	x				aux	
<i>Mercurialis annua</i>		neo	*		x	x		cwj	
<i>Mercurialis perennis</i>		nat	*		x		x	ckw	
<i>Meum athamanticum</i>		nat	V	x				auy	
<i>Milium effusum</i>		nat	*				x	cwl	
<i>Mimulus guttatus</i>		neo	*					auz	
<i>Mimulus moschatus</i>		neo	*					cwm	
<i>Mimuartia verna ssp. hercynica</i>		nat	3					avb	
<i>Misopates orontium</i>		arc	2					avd	
<i>Moehringia trinervia</i>		nat	*		x		x	cwn	
<i>Molinia caerulea</i>		nat	*	x				cwo	
<i>Moneses uniflora</i>		nat	1	x			x	avf	
<i>Monotropa hypopitys agg.</i>		nat	3	x				avg	cwp, cwq
<i>Montia fontana</i>		nat	3					avh	cwi, avi, cws, cwt
<i>Mycelis muralis</i>		nat	*					cxw	
<i>Myosotis arvensis</i>		arc	*					cwv	
<i>Myosotis discolor</i>		nat	V	x				avj	
<i>Myosotis laxa</i>		nat	*		x			avk	
<i>Myosotis nemorosa</i>		nat	*					avl	
<i>Myosotis ramosissima</i>		arc	*	x				avm	
<i>Myosotis scorpioides</i>		nat	*					cwz	
<i>Myosotis sparsiflora</i>		nat	R		x			avn	
<i>Myosotis stricta</i>		arc	V	x				cxv	
<i>Myosotis sylvatica</i>		nat	*		x		x	cxv	
<i>Myosurus minimus</i>		arc	*					avo	
<i>Myrica gale</i>		nat	3	x				avp	
<i>Myriophyllum alterniflorum</i>		nat	3	x				avq	
<i>Myriophyllum spicatum</i>		nat	*		x			avr	
<i>Myriophyllum verticillatum</i>		nat	*		x			avs	
<i>Myrrhis odorata</i>		neo	*		x	x		cxv	
<i>Nardus stricta</i>		nat	V	x				cxv	
<i>Narthecium ossifragum</i>		nat	3	x				avu	
<i>Nasturtium officinale agg.</i>		nat	*		x			edb	cxg, 191
<i>Neottia nidus-avis</i>		nat	*				x	avv	
<i>Nepeta cataria</i>		arc	2		x	x		avw	
<i>Neslia paniculata</i>		arc	1					avx	
<i>Nonea pulla</i>		nat	1	x				avz	
<i>Nuphar lutea</i>		nat	*					awa	
<i>Nuphar x spenneriana</i>		nat	R					cxr	
<i>Nymphaea alba</i>		nat	V					awc	
<i>Nymphaea candida</i>		nat	3					awd	
<i>Nymphoides peltata</i>		nat	2		x			awe	
<i>Odontites litoralis</i>	x	nat	1					awf	
<i>Odontites vernus</i>		nat	3					cxk	
<i>Odontites vulgaris</i>		nat	*					cxl	
<i>Oenanthe aquatica</i>		nat	*					cxm	
<i>Oenanthe coniodes</i>	x	nat	1		x			awh	
<i>Oenanthe fistulosa</i>		nat	3					awi	
<i>Oenanthe lachenalii</i>		nat	3		x			awj	
<i>Oenothera biennis</i>		neo	*					cxo	
<i>Oenothera oakesiana</i>		neo	*					cxn	
<i>Omphalodes scorpioides</i>		nat	1		x		x	awl	
<i>Onobrychis vicifolia</i>		neo	*	x				cxv	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Ononis spinosa</i> agg.		nat	*					awn	awm, awo
<i>Onopordum acanthium</i>		arc	*		x	x		awp	
<i>Ophioglossum vulgatum</i>		nat	3	x				awq	
<i>Ophrys apifera</i>		nat	3	x				awr	
<i>Ophrys insectifera</i>		nat	3	x				aws	
<i>Orchis mascula</i>		nat	3					awu	
<i>Orchis militaris</i>		nat	2	x				awv	
<i>Orchis purpurea</i>		nat	3	x				awy	
<i>Orchis tridentata</i>		nat	2	x				awz	
<i>Oreopteris limbosperma</i>		nat	*				x	bgu	
<i>Origanum vulgare</i>		nat	*	x				axb	
<i>Ornithogalum nutans</i> agg.		neo	*		x	x		axd	
<i>Ornithogalum umbellatum</i>		arc	*		x	x		axe	
<i>Ornithopus perpusillus</i>		nat	*	x				cxw	
<i>Ornithopus sativus</i>		neo	*					cxx	
<i>Orobanche caryophyllacea</i>		nat	1	x				axh	
<i>Orobanche elatior</i>		nat	1	x				axi	
<i>Orobanche lutea</i>		nat	1					axj	
<i>Orobanche picridis</i>		nat	1					axm	
<i>Orobanche purpurea</i>		nat	1	x				axn	
<i>Orobanche rapum-genistae</i>		nat	2	x				axp	
<i>Orobanche reticulata</i>		nat	3	x				axl	
<i>Orthilia secunda</i>		nat	3	x			x	axq	
<i>Osmunda regalis</i>		nat	3				x	axr	
<i>Oxalis acetosella</i>		nat	*				x	cxz	
<i>Oxalis corniculata</i>		neo	*			x		cya	
<i>Oxalis stricta</i>		neo	*		x			cyc	
<i>Panicum riparium</i>		n/a	*					dsf	
<i>Papaver argemone</i>		arc	*					cyh	
<i>Papaver dubium</i>		arc	*					ede	cyi, cyj
<i>Papaver rhoeas</i>		arc	*					axs	
<i>Parapholis strigosa</i>		nat	*					cyn	
<i>Parietaria judaica</i>		neo	*		x	x		axt	
<i>Parietaria officinalis</i>		arc	3		x	x		axu	
<i>Paris quadrifolia</i>		nat	V		x		x	axv	
<i>Parnassia palustris</i>		nat	2	x				axw	
<i>Pastinaca sativa</i>		nat	*					cyr	
<i>Pedicularis palustris</i>		nat	2	x				axx	
<i>Pedicularis sylvatica</i>		nat	2	x				axy	
<i>Peplis portula</i>		nat	V	x				axz	
<i>Persicaria amphibia</i>		nat	*					daf	
<i>Persicaria hydropiper</i>		nat	*		x			dak	
<i>Persicaria lapathifolia</i>		nat	*					edl	dal, dam
<i>Persicaria maculosa</i>		nat	*		x			dap	
<i>Persicaria minor</i>		nat	*		x			dao	
<i>Persicaria mitis</i>		n/a	*		x			azh	
<i>Petasites albus</i>		nat	*					cys	
<i>Petasites hybridus</i>		nat	*		x			cyt	
<i>Petasites spurius</i>		nat	2					aya	
<i>Petrohragia prolifera</i>		nat	2	x				ayb	
<i>Peucedanum cervaria</i>		nat	2	x				ayc	
<i>Peucedanum officinale</i>	x	nat	1	x				ayd	
<i>Peucedanum oreoselinum</i>		nat	2	x				aye	
<i>Peucedanum ostruthium</i>		neo	R		x			ayf	
<i>Peucedanum palustre</i>		nat	*					ayg	
<i>Phalaris arundinacea</i>		nat	*		x			cyv	
<i>Phegopteris connectilis</i>		nat	*				x	bgw	
<i>Phleum arenarium</i>	x	nat	V	x				ayh	
<i>Phleum bertolonii</i>		nat	*					cyx	
<i>Phleum paniculatum</i>	x	nat	1					ayi	
<i>Phleum phleoides</i>		nat	2	x				ayj	
<i>Phleum pratense</i>		nat	*		x			cyy	
<i>Phragmites australis</i>		nat	*		x			cyz	
<i>Physalis alkekengi</i>		nat	*		x			ayl	
<i>Phyteuma nigrum</i>		nat	V				x	aym	
<i>Phyteuma orbiculare</i>		nat	2	x				ayn	
<i>Phyteuma spicatum</i>		nat	*				x	ayo	cza, czb
<i>Picea abies</i>		nat	*					czc	
<i>Picris hieracioides</i>		nat	*					cze	
<i>Pilularia globulifera</i>		nat	2	x				ayp	
<i>Pimpinella major</i>		nat	*					czf	
<i>Pimpinella nigra</i>		nat	n/a	x				402	
<i>Pimpinella saxifraga</i>		nat	*	x				czg	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Pinguicula vulgaris</i>		nat	2	x				ayq	
<i>Pinus sylvestris</i>		nat	*					czi	
<i>Plantago coronopus</i>		nat	*					ayr	
<i>Plantago lanceolata</i>		arc	*					czj	
<i>Plantago major</i>		nat	*					edi	czk, czl, ayt
<i>Plantago maritima</i>		nat	*					ayu	
<i>Plantago media</i>		nat	*	x				czm	
<i>Platanthera bifolia</i>		nat	2					ayv	
<i>Platanthera chlorantha</i>		nat	3				x	ayw	
<i>Poa angustifolia</i>		nat	*	x				czn	
<i>Poa annua</i>		nat	*		x			czo	
<i>Poa bulbosa</i>		nat	3	x				ayx	
<i>Poa chaixii</i>		nat	*				x	czp	
<i>Poa compressa</i>		nat	*	x				czq	
<i>Poa humilis</i>		nat	*	x				czu	
<i>Poa nemoralis</i>		nat	*				x	czr	
<i>Poa palustris</i>		nat	*		x			czs	
<i>Poa pratensis</i>		nat	*					czt	
<i>Poa remota</i>		nat	3		x		x	ayy	
<i>Poa supina</i>		nat	*		x			czv	
<i>Poa trivialis</i>		nat	*		x			czw	
<i>Polemonium caeruleum</i>	x	nat	R					czx	
<i>Polygala amara</i> agg.		nat	2	x				azc	daa, dab
<i>Polygala comosa</i>		nat	V	x				dac	
<i>Polygala serpyllifolia</i>		nat	2	x				azd	
<i>Polygala vulgaris</i>		nat	3	x				aze	cyk, cyq
<i>Polygonatum multiflorum</i>		nat	*				x	dad	
<i>Polygonatum odoratum</i>		nat	2	x				azf	
<i>Polygonatum verticillatum</i>		nat	*				x	dae	
<i>Polygonum aviculare</i> agg.		nat	*					dag	144
<i>Polypodium vulgare</i> agg.		nat	*	x				edm	dar, 276
<i>Polystichum aculeatum</i>		nat	3		x		x	azi	
<i>Polystichum lonchitis</i>		nat	R	x				azj	
<i>Populus alba</i>		neo	*					dat	
<i>Populus nigra</i>		nat	3		x			azk	
<i>Populus tremula</i>		nat	*					dau	
<i>Populus x canescens</i>		neo	*					258	
<i>Portulaca oleracea</i>		neo	*		x	x		dav	
<i>Potamogeton acutifolius</i>		nat	3					azl	
<i>Potamogeton alpinus</i>		nat	V					azm	
<i>Potamogeton coloratus</i>		nat	2		x			azo	
<i>Potamogeton compressus</i>		nat	3					azp	
<i>Potamogeton crispus</i>		nat	*					dax	
<i>Potamogeton friesii</i>		nat	3					azr	
<i>Potamogeton gramineus</i>		nat	2					azs	
<i>Potamogeton lucens</i>		nat	3		x			azt	
<i>Potamogeton natans</i>		nat	*					day	
<i>Potamogeton nodosus</i>		nat	3					azu	
<i>Potamogeton obtusifolius</i>		nat	3					azv	
<i>Potamogeton pectinatus</i>		nat	*		x			daz	
<i>Potamogeton perfoliatus</i>		nat	3					azw	
<i>Potamogeton polygonifolius</i>		nat	3	x				azx	
<i>Potamogeton praelongus</i>		nat	1					azy	
<i>Potamogeton pusillus</i> agg.		nat	*					edo	daw
<i>Potamogeton trichoides</i>		nat	*					dbb	
<i>Potamogeton x angustifolius</i>		nat	1					azn	
<i>Potamogeton x nitens</i>		nat	1					dbe	
<i>Potamogeton x spathulatus</i>		nat	G					dbg	
<i>Potamogeton x undulatus</i>		n/a	R					dbh	
<i>Potentilla alba</i>		nat	1					baa	
<i>Potentilla anglica</i>		nat	V					bab	
<i>Potentilla anserina</i>		nat	*		x			dbi	
<i>Potentilla argentea</i>		nat	*	x				dbj	
<i>Potentilla erecta</i>		nat	*	x				dbk	
<i>Potentilla heptaphylla</i>		nat	2	x				bae	
<i>Potentilla incana</i>		nat	1	x				bac	
<i>Potentilla intermedia</i>		neo	*			x		dbm	
<i>Potentilla neummanniana</i>		nat	V	x				baf	
<i>Potentilla norvegica</i>		neo	*					dbn	
<i>Potentilla palustris</i>		nat	V	x				bag	
<i>Potentilla recta</i>		neo	*	x				dbo	
<i>Potentilla reptans</i>		nat	*					dbp	
<i>Potentilla sterilis</i>		nat	*				x	bah	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Potentilla supina</i>		nat	3		x			bai	
<i>Primula elatior</i>		nat	*		x		x	baj	
<i>Primula veris</i>		nat	V	x				bak	
<i>Primula vulgaris</i>		nat	1				x	bal	
<i>Prunella grandiflora</i>		nat	3	x				bam	
<i>Prunella laciniata</i>		nat	2	x				ban	
<i>Prunella vulgaris</i>		nat	*					dbr	
<i>Prunus avium</i>		nat	*					dbb	
<i>Prunus padus</i>		nat	*					dbx	
<i>Prunus serotina</i>		neo	*					dby	
<i>Prunus spinosa</i>		nat	*					dbz	
<i>Pseudognaphalium luteoalbum</i>		nat	2	x				apf	
<i>Pseudolysimachion longifolium</i>		nat	3					biw	
<i>Pseudolysimachion spicatum</i>		nat	2	x				bjc	
<i>Pseudorchis albida</i>		nat	1	x				bao	
<i>Psyllium arenarium</i>		neo	*					ays	
<i>Pteridium aquilinum</i>		nat	*	x			x	dcb	
<i>Puccinellia capillaris</i>	x	nat	R					dcc	
<i>Puccinellia distans</i>		nat	*					dcd	
<i>Puccinellia limosa</i>		nat	R					89	
<i>Puccinellia maritima</i>		nat	*					dce	
<i>Pulicaria dysenterica</i>		nat	3					baq	
<i>Pulicaria vulgaris</i>		nat	3		x			bar	
<i>Pulmonaria obscura</i>		nat	*		x		x	dcf	
<i>Pulmonaria officinalis</i>		nat	*				x	bas	
<i>Pulsatilla pratensis</i>		nat	2	x				bau	
<i>Pulsatilla vulgaris</i>		nat	2	x				baw	
<i>Pyrola chlorantha</i>		nat	1	x			x	bax	
<i>Pyrola minor</i>		nat	3	x			x	baz	
<i>Pyrola rotundifolia</i>		nat	3	x			x	bba	
<i>Pyrus pyraeaster</i>		nat	3					bbb	
<i>Quercus petraea</i>		nat	*					dch	
<i>Quercus robur</i>		nat	*					dci	
<i>Quercus x rosacea</i>		nat	*					dsu	
<i>Radiola linoides</i>		nat	3	x				bbc	
<i>Ranunculus acris</i>		nat	*					dcl	
<i>Ranunculus aquatilis</i> agg.		nat	*					bbd	bbe, bbi, bbj, bbk, bbn, bbg, bbo, dco, bbx
<i>Ranunculus arvensis</i>		arc	2					bbf	
<i>Ranunculus auricomus</i> agg.		nat	*					dck	
<i>Ranunculus bulbosus</i>		nat	*	x				bbh	
<i>Ranunculus ficaria</i> ssp. <i>bulbilifer</i>		nat	*		x			dcl	
<i>Ranunculus flammula</i>		nat	*	x				dcm	
<i>Ranunculus lanuginosus</i>		nat	*		x		x	dcm	
<i>Ranunculus lingua</i>		nat	3		x			bbi	
<i>Ranunculus platanifolius</i>		nat	3		x			bbp	
<i>Ranunculus polyanthemus</i> agg.		nat	3					bbs	bbm, bbq, bbr, bbw, bbt
<i>Ranunculus repens</i>		nat	*		x			dcp	
<i>Ranunculus sardous</i>		nat	3		x			bbv	
<i>Ranunculus sceleratus</i>		nat	*		x			dcq	
<i>Raphanus raphanistrum</i>		arc	3					dcr	
<i>Reseda lutea</i>		arc	*					dcs	
<i>Reseda luteola</i>		arc	*					dcs	
<i>Rhamnus cathartica</i>		nat	*					bbz	
<i>Rhinanthus alectorolophus</i>		nat	3					bca	
<i>Rhinanthus angustifolius</i> ssp. <i>grandiflorus</i>		nat	V	x				bcd	
<i>Rhinanthus glacialis</i>		nat	1	x				bcb	
<i>Rhinanthus minor</i>		nat	V	x				bcc	
<i>Rhynchospora alba</i>		nat	3	x				bce	
<i>Rhynchospora fusca</i>		nat	2	x				bcd	
<i>Ribes alpinum</i>		nat	*		x		x	dcs	
<i>Ribes nigrum</i>		nat	*				x	dcs	
<i>Ribes rubrum</i> agg.		nat	*					edw	
<i>Ribes uva-crispa</i>		nat	*			x		dda	
<i>Robinia pseudoacacia</i>		neo	*		x			ddb	
<i>Rorippa amphibia</i>		nat	*		x			ddc	
<i>Rorippa anceps</i>		nat	*		x			bcd	
<i>Rorippa austriaca</i>		nat	R		x			bci	
<i>Rorippa palustris</i>		nat	*		x			ddd	
<i>Rorippa sylvestris</i>		nat	*					dde	
<i>Rosa agrestis</i>		nat	2	x				bcj	
<i>Rosa arvensis</i>		nat	2				x	bck	
<i>Rosa caesia</i> agg.		nat	n/a					bcl	ddf
<i>Rosa canina</i>		nat	*					ddg	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Rosa corymbifera</i>		nat	*					ddh	
<i>Rosa dumalis</i>		nat	3					dds	bcq
<i>Rosa elliptica</i>		nat	3	x				bcm	
<i>Rosa inodora</i>		nat	G					87	
<i>Rosa micrantha</i>		nat	3	x				ddj	
<i>Rosa rubiginosa</i>		nat	*	x				ddk	
<i>Rosa rugosa</i>		neo	*					ddl	
<i>Rosa spinosissima</i>	x	nat	3	x				bco	
<i>Rosa subcanina</i>		nat	*	x				ddo	
<i>Rosa subcollina</i>		nat	3	x				ddp	
<i>Rosa tomentella</i>		nat	3					bcn	
<i>Rosa tomentosa</i> agg.		nat	3					bcp	ddm, ddn, ddq, ddr
<i>Rubus armeniacus</i>		neo	*			x		dea	
<i>Rubus caesius</i>		nat	*		x			deg	
<i>Rubus chamaemorus</i>		nat	2	x				bcr	
<i>Rubus fruticosus</i> agg.		nat	*					edy	
<i>Rubus idaeus</i>		nat	*				x	dgi	
<i>Rubus laciniatus</i>		neo	*					dgo	
<i>Rubus saxatilis</i>		nat	2				x	bcs	
<i>Rumex acetosa</i>		nat	*					djt	
<i>Rumex acetosella</i>		nat	*	x				dju	
<i>Rumex aquaticus</i>		nat	V		x			bet	
<i>Rumex conglomeratus</i>		nat	*		x			djv	
<i>Rumex crispus</i>		nat	*					djw	
<i>Rumex hydrolapathum</i>		nat	*		x			djx	
<i>Rumex maritimus</i>		nat	*		x			bcu	
<i>Rumex obtusifolius</i>		nat	*		x			eeb	
<i>Rumex palustris</i>		nat	*		x			bcv	
<i>Rumex salicifolius</i>	x	neo	*		x			dke	
<i>Rumex sanguineus</i>		nat	*		x		x	dka	
<i>Rumex stenophyllus</i>		neo	*					dkc	
<i>Rumex thyrsiflorus</i>		nat	*					dkd	
<i>Rumex x pratensis</i>		n/a	*					far	
<i>Ruppia cirrhosa</i>	x	nat	2					dkf	
<i>Ruppia maritima</i>	x	nat	2					bcw	
<i>Sagina apetala</i> agg.		arc	*					eed	dkh
<i>Sagina maritima</i>		nat	*	x				dki	
<i>Sagina nodosa</i>		nat	3					bcx	
<i>Sagina procumbens</i>		nat	*			x		dkj	
<i>Sagittaria latifolia</i>		neo	*					dkk	
<i>Sagittaria sagittifolia</i>		nat	*					bcz	
<i>Salicornia dolichostachya</i> s.l.		nat	n/a	x				eef	
<i>Salicornia europaea</i>		nat	*					eee	bda, dkn
<i>Salicornia fragilis</i>	x	nat	n/a					dko	
<i>Salicornia procumbens</i>		nat	*	x				dkl	
<i>Salicornia stricta</i>		nat	*					dkm	
<i>Salix alba</i>		nat	*		x			eeg	dkq
<i>Salix aurita</i>		nat	*	x				dkr	
<i>Salix caprea</i>		nat	*		x			dks	
<i>Salix cinerea</i>		nat	*					dkt	
<i>Salix fragilis</i> agg.		nat	*					eeh	dkv, dle
<i>Salix pentandra</i>		nat	3					dkw	
<i>Salix purpurea</i>		nat	*					eei	dky
<i>Salix repens</i>		nat	*					eej	dla, bdd
<i>Salix triandra</i>		nat	*					eek	dlb
<i>Salix viminalis</i>		nat	*					dld	
<i>Salix x alopecuroides</i>		nat	*					101	
<i>Salix x ambigua</i>		nat	*					drc	
<i>Salix x holosericea</i>		nat	*					drb	
<i>Salix x meyeriana</i>		nat	*					dqr	
<i>Salix x mollissima</i>		nat	*					dqi	
<i>Salix x multinervis</i>		nat	*					dkx	
<i>Salix x reichardtii</i>		nat	*					dqh	
<i>Salix x rubra</i>		nat	*					dqf	
<i>Salix x smithiana</i>		nat	*					dli	
<i>Salsola kali</i>		nat	*					bdf	dlg, dlh
<i>Salvia glutinosa</i>		neo	*		x			dli	
<i>Salvia nemorosa</i>		neo	*					dli	
<i>Salvia pratensis</i>		nat	3					bdg	
<i>Salvia verticillata</i>		neo	*				x	bdh	
<i>Sambucus ebulus</i>		neo	*		x	x	x	dli	
<i>Sambucus nigra</i>		nat	*		x			dli	14
<i>Sambucus racemosa</i>		nat	*		x			dli	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Samolus valerandi</i>		nat	2					bdi	
<i>Sanguisorba minor</i>		nat	*	x				eel	bdj
<i>Sanguisorba officinalis</i>		nat	3					bdk	
<i>Sanicula europaea</i>		nat	*					bdl	
<i>Saponaria officinalis</i>		nat	*			x		dip	
<i>Saxifraga granulata</i>		nat	3	x				bdm	
<i>Saxifraga tridactylites</i>		nat	*	x				bdo	
<i>Scabiosa canescens</i>		nat	2	x				bdp	
<i>Scabiosa columbaria</i>		nat	*	x				bdq	
<i>Scandix pecten-veneris</i>		arc	2					bds	
<i>Scheuchzeria palustris</i>		nat	2	x				bdt	
<i>Schoenoplectus lacustris</i>		nat	*					dlt	
<i>Schoenoplectus pungens</i>		nat	1		x			bdu	
<i>Schoenoplectus tabernaemontani</i>		nat	*					bdw	
<i>Schoenoplectus triquetet</i>		nat	3		x			bdx	
<i>Schoenoplectus x carinatus</i>		nat	3		x			bdy	
<i>Schoenus nigricans</i>	x	nat	3	x				bdz	
<i>Scirpus sylvaticus</i>		nat	*					dlu	
<i>Scleranthus annuus</i> agg.		arc	*					een	een
<i>Scleranthus perennis</i>		nat	3	x				bea	
<i>Scorzonera hispanica</i>	x	nat	2	x				beb	
<i>Scorzonera humilis</i>		nat	2	x				bec	
<i>Scorzonera laciniata</i>		nat	2					ayz	
<i>Scrophularia nodosa</i>		nat	*		x			dly	
<i>Scrophularia umbrosa</i>		nat	*		x			bee	
<i>Scrophularia vernalis</i>		neo	*		x	x		dlz	
<i>Scutellaria galericulata</i>		nat	*					dma	
<i>Scutellaria hastifolia</i>		nat	2					bef	
<i>Scutellaria minor</i>		nat	2	x				beg	
<i>Securigera varia</i>		neo	*	x				ajo	
<i>Sedum acre</i>		nat	*	x				dmb	
<i>Sedum album</i>		nat	*	x				beh	
<i>Sedum rupestre</i>		nat	V	x				bei	
<i>Sedum sexangulare</i>		nat	*	x				bej	
<i>Sedum telephium</i> agg.		nat	*					eoo	dmd, dme
<i>Selinum carvifolia</i>		nat	3	x				bel	
<i>Sempervivum tectorum</i>		nat	*					bem	
<i>Senecio aquaticus</i>		nat	3					ben	
<i>Senecio erraticus</i> ssp. <i>barbareifolius</i>		nat	3					bep	
<i>Senecio erucifolius</i>		nat	*					dmg	
<i>Senecio hercynicus</i>		nat	*		x		x	bes	
<i>Senecio inaequidens</i>		neo	*	x		x		dmh	
<i>Senecio jacobaea</i> ssp. <i>dunensis</i>	x	nat	*					dmi	
<i>Senecio jacobaea</i> ssp. <i>jacobaea</i>		nat	*					dmj	
<i>Senecio ovatus</i>		nat	*		x		x	dmk	
<i>Senecio paludosus</i>		nat	2					bet	
<i>Senecio sarracenicus</i>		nat	*		x			beq	
<i>Senecio sylvaticus</i>		nat	*		x		x	dml	
<i>Senecio vernalis</i>		neo	*					dmm	
<i>Senecio viscosus</i>		nat	*					dmn	
<i>Senecio vulgaris</i>		nat	*		x			dmo	
<i>Serratula tinctoria</i>		nat	2	x				beu	
<i>Seseli annuum</i>		nat	1	x				bev	
<i>Seseli libanotis</i>		nat	2	x				bew	
<i>Seseli montanum</i>		nat	1					dmp	
<i>Sesleria albicans</i>		nat	V	x				bex	
<i>Setaria pumila</i>		arc	V					bey	
<i>Setaria viridis</i>		arc	*		x			dmt	
<i>Sherardia arvensis</i>		arc	3					bez	
<i>Silaum silaus</i>		nat	2	x				bfa	
<i>Silene conica</i>	x	neo	*	x				dmw	
<i>Silene dioica</i>		nat	*		x			dmx	
<i>Silene flos-cuculi</i>		nat	*					cuy	
<i>Silene latifolia</i> ssp. <i>alba</i>		nat	*		x			dmz	
<i>Silene noctiflora</i>		arc	3					bfc	
<i>Silene nutans</i>		nat	V	x				bfd	
<i>Silene otites</i>		nat	3	x				bfe	
<i>Silene viscaria</i>		nat	1	x				aud	
<i>Silene vulgaris</i>		nat	*					bff	
<i>Silene x hampeana</i>		n/a	*					dqw	
<i>Sinapis arvensis</i>		arc	*					dnd	
<i>Sisymbrium altissimum</i>		neo	*					dne	
<i>Sisymbrium austriacum</i>		nat	R		x			bfg	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Sisymbrium loeselii</i>		neo	*					dnq	
<i>Sisymbrium officinale</i>		arc	*		x			dnh	
<i>Sisymbrium strictissimum</i>		nat	R		x			bfi	
<i>Sium latifolium</i>		nat	*		x			bfi	
<i>Solanum dulcamara</i>		nat	*		x			dnk	
<i>Solanum nigrum</i>		arc	*		x			eeq	dnm
<i>Solidago canadensis</i>		neo	*					dns	
<i>Solidago gigantea</i>		neo	*		x			dnt	
<i>Solidago virgaurea</i>		nat	*					dnv	
<i>Sonchus arvensis</i>		nat	*					eer	dnw, bfk
<i>Sonchus asper</i>		nat	*		x			dnx	
<i>Sonchus oleraceus</i>		nat	*		x			dny	
<i>Sonchus palustris</i>		nat	V		x			bfi	
<i>Sorbus aucuparia</i>		nat	*					dnz	
<i>Sorbus torminalis</i>		nat	*					bfo	
<i>Sparganium angustifolium</i>		nat	2	x				bfp	
<i>Sparganium emersum</i>		nat	*		x			doa	
<i>Sparganium erectum</i>		nat	*		x			ees	doc
<i>Sparganium natans</i>		nat	2	x				bfq	
<i>Spartina anglica</i>		neo	*	x				dod	
<i>Spergula arvensis</i>		arc	*					doe	
<i>Spergula morisonii</i>		nat	*	x				dof	
<i>Spergularia echinosperma</i>		nat	*		x			bfr	
<i>Spergularia media</i>		nat	*					dog	
<i>Spergularia rubra</i>		arc	*					doh	
<i>Spergularia salina</i>		nat	*					bfs	
<i>Spiraea alba</i>		neo	*					dpf	
<i>Spiranthes spiralis</i>		nat	1	x				bfu	
<i>Spirodela polyrrhiza</i>		nat	*					dok	
<i>Stachys alpina</i>		nat	3		x		x	bfv	
<i>Stachys annua</i>		arc	1					bfw	
<i>Stachys arvensis</i>		arc	3					bfx	
<i>Stachys germanica</i>		nat	2					bfy	
<i>Stachys palustris</i>		nat	*					dol	
<i>Stachys recta</i>		nat	2	x				bfz	
<i>Stachys sylvatica</i>		nat	*		x		x	dom	
<i>Stachys x ambigua</i>		nat	*					103	
<i>Stellaria alsine</i>		nat	*					dot	
<i>Stellaria aquatica</i>		nat	*		x			dou	
<i>Stellaria graminea</i>		nat	*	x				doo	
<i>Stellaria holostea</i>		nat	*				x	dop	
<i>Stellaria media</i>		arc	*					doq	
<i>Stellaria neglecta</i>		nat	*				x	dor	
<i>Stellaria nemorum</i>		nat	*		x			bgb	
<i>Stellaria pallida</i>		arc	*					dos	
<i>Stellaria palustris</i>		nat	V	x				bgc	
<i>Stipa capillata</i>	x	nat	R	x				bgd	
<i>Stratiotes aloides</i>		nat	3					bge	
<i>Suaeda maritima</i>		nat	*		x			dou	
<i>Succisa pratensis</i>		nat	3	x				dov	
<i>Symphytum officinale</i>		nat	*		x			doy	
<i>Symphytum x uplandicum</i>		neo	*					doz	
<i>Tanacetum corymbosum</i>		nat	3				x	bgf	
<i>Tanacetum vulgare</i>		arc	*			x		dpc	
<i>Taraxacum acervatulum</i>		nat	n/a					104	
<i>Taraxacum acutifrons</i>		nat	n/a					dpe	
<i>Taraxacum adiantifrons</i>		nat	n/a					dqn	
<i>Taraxacum aequilobum</i>		nat	n/a					106	
<i>Taraxacum alatum</i>		nat	n/a					dph	
<i>Taraxacum amphibolum</i>		nat	n/a					107	
<i>Taraxacum amplum</i>		nat	n/a					dpi	
<i>Taraxacum ancistrolobum</i>		nat	n/a					dpj	
<i>Taraxacum angustisquamum</i>		nat	n/a					108	
<i>Taraxacum atactum</i>		nat	n/a					dpk	
<i>Taraxacum atricapillum</i>	x	nat	n/a					110	
<i>Taraxacum baecifforme</i>		nat	n/a					dpn	
<i>Taraxacum borgvallii</i>		nat	n/a					dpp	
<i>Taraxacum cacuminatum</i>		nat	n/a					111	
<i>Taraxacum calochroum</i>	x	nat	n/a					112	
<i>Taraxacum caloschistum</i>		nat	n/a					dpu	
<i>Taraxacum canoviride</i>		nat	n/a					113	
<i>Taraxacum celticum agg.</i>		nat	V					eex	dpr, dsa, 26, 283, drg, 181, 182, dqj
<i>Taraxacum chlorodes</i>	x	nat	n/a					114	

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Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Taraxacum christiansenii</i>	x	nat	n/a					115	
<i>Taraxacum contulium</i>		nat	n/a					117	
<i>Taraxacum contractum</i>		nat	n/a					119	
<i>Taraxacum copidophyllum</i>		nat	n/a					120	
<i>Taraxacum cordatum</i>		nat	n/a					dpw	
<i>Taraxacum croceiflorum</i>	x	nat	n/a					dpz	
<i>Taraxacum cyanolepis</i>		nat	n/a					dqa	
<i>Taraxacum dilaceratum</i>		nat	n/a					dqb	
<i>Taraxacum dilatatum</i>	x	nat	n/a					121	
<i>Taraxacum edytomum</i>		nat	n/a					140	
<i>Taraxacum ekmannii</i>		nat	n/a					dqd	
<i>Taraxacum exsertiforme</i>		nat	n/a					123	
<i>Taraxacum exsertum</i>		nat	n/a					124	
<i>Taraxacum fagerstroemii</i>		nat	n/a					125	
<i>Taraxacum fasciatum</i>		nat	n/a					dqg	
<i>Taraxacum fulgidum</i>		nat	n/a					127	
<i>Taraxacum fusciflorum</i>		nat	n/a					57	
<i>Taraxacum gelertiiforme</i>		nat	n/a					128	
<i>Taraxacum gesticulans</i>		nat	n/a					129	
<i>Taraxacum hamatiforme</i>		nat	n/a					dqk	
<i>Taraxacum hamatululum</i>		nat	n/a					58	
<i>Taraxacum hamatum</i>		nat	n/a					130	
<i>Taraxacum hamatum agg.</i>		nat	*					dql	
<i>Taraxacum hamiferum</i>	x	nat	n/a					dqm	
<i>Taraxacum hepaticum</i>		nat	n/a					131	
<i>Taraxacum huelphersianum</i>		nat	n/a					dqo	
<i>Taraxacum inarmatum</i>		nat	n/a					132	
<i>Taraxacum ingens</i>		nat	n/a					dqq	
<i>Taraxacum inops</i>		nat	n/a					133	
<i>Taraxacum interveniens</i>		nat	n/a					134	
<i>Taraxacum intumescens</i>		nat	n/a					135	
<i>Taraxacum kernianum</i>		nat	n/a					dqs	
<i>Taraxacum lacinosifrons</i>		nat	n/a					dqt	
<i>Taraxacum lacinosum</i>	x	nat	n/a					dqu	
<i>Taraxacum lacinulatum</i>		nat	n/a					136	
<i>Taraxacum laevigatum agg.</i>		nat	*	x				bgi	dpq, 116, 122, 56, 126, dqv, dsg, drw, 34, dsd, dsr, dss
<i>Taraxacum lamprophyllum</i>		nat	n/a					dqx	
<i>Taraxacum latens</i>		nat	n/a					137	
<i>Taraxacum laticordatum</i>		nat	n/a					dqy	
<i>Taraxacum latisectum</i>	x	nat	n/a					138	
<i>Taraxacum latissimum</i>		nat	n/a					139	
<i>Taraxacum leptodon</i>		nat	n/a					dqz	
<i>Taraxacum lingulatum</i>		nat	n/a					142	
<i>Taraxacum lojoense</i>		nat	n/a					143	
<i>Taraxacum lucidum</i>		nat	n/a					drd	
<i>Taraxacum macranthoides</i>	x	nat	n/a					145	
<i>Taraxacum maculatum</i>		nat	n/a					147	
<i>Taraxacum melanostigma</i>		nat	n/a					148	
<i>Taraxacum melanthoides</i>		nat	n/a					dre	
<i>Taraxacum mimulum</i>		nat	n/a					dri	
<i>Taraxacum necessarium</i>		nat	n/a					149	
<i>Taraxacum nitidum</i>		nat	n/a					150	
<i>Taraxacum obliquilobum</i>		nat	n/a					dri	
<i>Taraxacum obliquum</i>	x	nat	R					dri	
<i>Taraxacum oblongatum</i>		nat	n/a					dri	
<i>Taraxacum obtusifrons</i>		nat	n/a					drk	
<i>Taraxacum ochrochlorum</i>		nat	n/a					151	
<i>Taraxacum officinale agg.</i>		nat	*		x			eey	
<i>Taraxacum ostenfeldii</i>		nat	n/a					dqc	
<i>Taraxacum oxyrhinum</i>		nat	n/a					152	
<i>Taraxacum pachymerum</i>	x	nat	n/a					153	
<i>Taraxacum pallidipes</i>		nat	n/a					drm	
<i>Taraxacum palustre agg.</i>		nat	3	x				bgj	dri, 59, dpo, 40, 27
<i>Taraxacum pannucium</i>		nat	n/a					dro	
<i>Taraxacum pannulatum</i>		nat	n/a					159	
<i>Taraxacum pectinatiforme</i>		nat	n/a					drq	
<i>Taraxacum piceatum</i>		nat	n/a					drr	
<i>Taraxacum piceipictum</i>		nat	n/a					160	
<i>Taraxacum planum</i>		nat	n/a					105	
<i>Taraxacum plicatifrons</i>		nat	n/a					161	
<i>Taraxacum polyodon</i>		nat	n/a					drt	
<i>Taraxacum porrigenis</i>		nat	n/a					162	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Taraxacum privum</i>		nat	n/a					dru	
<i>Taraxacum pruinaum</i>		nat	n/a					183	
<i>Taraxacum pseudoretroflexum</i>		nat	n/a					163	
<i>Taraxacum pulcherrimum</i>		nat	n/a					164	
<i>Taraxacum pulchrifolium</i>		nat	n/a					drx	
<i>Taraxacum pulverulentum</i>	x	nat	n/a					156	
<i>Taraxacum purpureum</i>		nat	n/a					drs	
<i>Taraxacum quadrans</i>		nat	n/a					drz	
<i>Taraxacum rhodopodum</i>		nat	n/a					165	
<i>Taraxacum rigens</i>		nat	n/a					166	
<i>Taraxacum sagittipetens</i>		nat	n/a					dsc	
<i>Taraxacum sellandii</i>		nat	n/a					157	
<i>Taraxacum semiglobosum</i>		nat	n/a					167	
<i>Taraxacum severum</i>		nat	n/a					168	
<i>Taraxacum sinuatum</i>		nat	n/a					dsh	
<i>Taraxacum stereodes</i>		nat	n/a					dsj	
<i>Taraxacum subcaescescens</i>		nat	n/a					169	
<i>Taraxacum subdahlstedtii</i>		nat	n/a					170	
<i>Taraxacum subditivum</i>		nat	n/a					184	
<i>Taraxacum subericinum</i>		nat	n/a					185	
<i>Taraxacum subhamatum</i>		nat	n/a					dsk	
<i>Taraxacum subhielphersianum</i>		nat	n/a					171	
<i>Taraxacum sublaeticolor</i>	x	nat	n/a					dsl	
<i>Taraxacum subleucopodum</i>		nat	n/a					172	
<i>Taraxacum subserratifrons</i>		nat	n/a					173	
<i>Taraxacum subundulatum</i>		nat	n/a					dso	
<i>Taraxacum subxanthostigma</i>	x	nat	n/a					174	
<i>Taraxacum tenebricans</i>		nat	n/a					dsq	
<i>Taraxacum tumentilobum</i>		nat	n/a					dsv	
<i>Taraxacum undulatiflorum</i>		nat	n/a					175	
<i>Taraxacum undulatifforme</i>	x	nat	n/a					176	
<i>Taraxacum undulatum</i>		nat	n/a					177	
<i>Taraxacum unguifrons</i>	x	nat	n/a					178	
<i>Taraxacum valens</i>		nat	n/a					154	
<i>Taraxacum vanum</i>		nat	n/a					179	
<i>Taraxacum violaceinervosum</i>		nat	n/a					158	
<i>Taxus baccata</i>		nat	3					bgk	
<i>Teesdalia nudicaulis</i>		nat	*	x				bgj	
<i>Tephrosia palustris</i>		nat	2		x			beo	
<i>Tetragonolobus maritimus</i>		nat	1	x				bgm	
<i>Teucrium botrys</i>		arc	2	x				bgn	
<i>Teucrium scordium</i>		nat	2					bgp	
<i>Teucrium scorodonia</i>		nat	*	x			x	bgq	
<i>Thalictrum flavum</i>		nat	3					bgr	
<i>Thalictrum lucidum</i>		nat	3	x				bgs	
<i>Thalictrum minus ssp. minus</i>		nat	2					bgt	
<i>Thalictrum simplex ssp. tenuifolium</i>		nat	R					bgv	
<i>Thelypteris palustris</i>		nat	3					bgx	
<i>Thesium ebracteatum</i>		nat	1	x				bgy	
<i>Thesium linophyllum</i>		nat	1	x				bgz	
<i>Thesium pyrenaicum</i>		nat	2	x				dta	
<i>Thlaspi arvense</i>		arc	*					dtb	
<i>Thlaspi caerulescens</i>		nat	1					bha	
<i>Thlaspi calaminare</i>	x	nat	1	x				bhb	
<i>Thlaspi perfoliatum</i>		nat	V	x				bhc	
<i>Thymus praecox</i>		nat	R	x				dte	
<i>Thymus pulegioides</i>		nat	*	x				dtf	
<i>Thymus serpyllum</i>		nat	3	x				dtg	
<i>Tilia cordata</i>		nat	*					bhe	
<i>Tilia platyphyllos</i>		nat	*		x			bhf	
<i>Torilis japonica</i>		nat	*		x		x	dth	
<i>Torilis nodosa</i>		neo	3					dti	
<i>Tragopogon dubius</i>		nat	*					dtj	
<i>Tragopogon pratensis ssp. minor</i>		nat	D					95	
<i>Tragopogon pratensis ssp. orientalis</i>		nat	R					bhg	260, bhh, bhi
<i>Tragopogon pratensis ssp. pratensis</i>		nat	*					dhk	
<i>Trichomanes speciosum</i>		nat	R				x	bhj	
<i>Trichophorum cespitosum</i>		nat	3					dtm	
<i>Trientalis europaea</i>		nat	*	x			x	bhk	
<i>Trifolium alpestre</i>		nat	2	x				dtl	
<i>Trifolium arvense</i>		nat	*	x				bhl	
<i>Trifolium aureum</i>		nat	3	x				bhm	
<i>Trifolium campestre</i>		nat	*	x				bhn	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Trifolium dubium</i>		nat	*					dto	
<i>Trifolium fragiferum</i>		nat	*		x			bhl	
<i>Trifolium hybridum</i>		neo	*					dtp	
<i>Trifolium medium</i>		nat	*	x				bhm	
<i>Trifolium montanum</i>		nat	3	x				bhn	
<i>Trifolium pratense</i>		nat	*					dts	
<i>Trifolium repens</i>		nat	*					dtl	
<i>Trifolium spadicum</i>	x	nat	2	x				bhq	
<i>Trifolium striatum</i>		nat	2	x				bhr	
<i>Triglochin maritimum</i>		nat	*					bhs	
<i>Triglochin palustre</i>		nat	3	x				bht	
<i>Tripleurospermum perforatum</i>		arc	*					dtw	
<i>Trisetum flavescens</i>		nat	*					dtx	
<i>Trollius europaeus</i>		nat	2					bhu	
<i>Tuberaria guttata</i>	x	nat	R	x				bhv	
<i>Tulipa sylvestris</i>		arc	3			x		bhw	
<i>Tussilago farfara</i>		nat	*					dtv	
<i>Typha angustifolia</i>		nat	*		x			dtz	
<i>Typha latifolia</i>		nat	*		x			dua	
<i>Ulex europaeus</i>		neo	3	x				bhy	
<i>Ulmus glabra</i>		nat	*		x			dub	
<i>Ulmus laevis</i>		nat	3		x			bhz	
<i>Ulmus minor</i>		nat	3					duc	
<i>Urtica dioica</i> ssp. <i>dioica</i>		nat	*					dud	
<i>Urtica urens</i>		arc	*		x	x		due	
<i>Utricularia australis</i>	x	nat	3	x				bia	
<i>Utricularia intermedia</i>		nat	1	x				bib	
<i>Utricularia minor</i>		nat	3	x				bic	
<i>Utricularia vulgaris</i> agg.		nat	3					bie	bif
<i>Vaccinium angustifolium</i> x <i>corymbosum</i>		neo	*					ea	
<i>Vaccinium macrocarpon</i>		neo	*	x				duf	
<i>Vaccinium myrtillus</i>		nat	*	x				dug	
<i>Vaccinium oxycoccos</i>		nat	3	x				bii	
<i>Vaccinium uliginosum</i>		nat	3	x				bij	
<i>Vaccinium vitis-idaea</i>		nat	*	x				bik	
<i>Vaccinium</i> x <i>intermedium</i>		nat	*					bih	
<i>Valeriana dioica</i>		nat	V	x				bil	
<i>Valeriana officinalis</i> agg.		nat	*					efc	bim, bin
<i>Valerianella carinata</i>		nat	*					bio	
<i>Valerianella dentata</i>		nat	3					bip	
<i>Valerianella locusta</i>		arc	*					duj	
<i>Valerianella ramosa</i>		nat	2					biq	
<i>Verbascum blattaria</i>		neo	R					duk	
<i>Verbascum densiflorum</i>		nat	*					dul	
<i>Verbascum lychnitis</i>		nat	*	x				bir	
<i>Verbascum nigrum</i>		nat	*		x			dum	
<i>Verbascum phlomoides</i>		nat	*					dun	
<i>Verbascum phoeniceum</i>		nat	1	x		x		bis	
<i>Verbascum thapsus</i>		nat	*		x			bit	
<i>Verbena officinalis</i>		arc	V		x	x		biu	
<i>Veronica agrestis</i>		arc	V		x			dup	
<i>Veronica anagallis-aquatica</i> agg.		nat	*					biv	duu
<i>Veronica arvensis</i>		arc	*					dus	
<i>Veronica beccabunga</i>		nat	*					dut	
<i>Veronica chamaedrys</i>		nat	*					duv	
<i>Veronica filiformis</i>		neo	*		x	x		duw	
<i>Veronica hederifolia</i>		nat	*		x			efd	duy
<i>Veronica montana</i>		nat	*				x	bix	
<i>Veronica officinalis</i>		nat	*					duz	
<i>Veronica opaca</i>		arc	3					biy	
<i>Veronica peregrina</i>		neo	*					dva	
<i>Veronica persica</i>		neo	*		x			dvb	
<i>Veronica polita</i>		arc	*		x			biz	
<i>Veronica praecox</i>		nat	2	x				bja	
<i>Veronica scutellata</i>		nat	V	x				bjb	
<i>Veronica serpyllifolia</i>		nat	*					dvc	
<i>Veronica teucrium</i>		nat	V	x				bjd	
<i>Veronica triphyllos</i>		arc	3					bje	
<i>Veronica verna</i>		nat	2	x				bjf	
<i>Viburnum opulus</i>		nat	*					dve	
<i>Vicia angustifolia</i> ssp. <i>angustifolia</i>		arc	*					dv1	
<i>Vicia angustifolia</i> ssp. <i>segetalis</i>		neo	*					256	
<i>Vicia cassubica</i>		nat	3	x			x	bjg	

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Table A.2 – Continued from previous page

Species	Edge	Natural.	RL NB	EIV N1-3	EIV N7-9	Urban	Forest	Code	Incl.
<i>Vicia cracca</i>		nat	*					dvf	
<i>Vicia dumetorum</i>		nat	3				x	bjh	
<i>Vicia hirsuta</i>		nat	*					dvh	
<i>Vicia lathyroides</i>		nat	3	x				bji	
<i>Vicia pisiformis</i>		nat	2	x			x	bjj	
<i>Vicia sativa</i> agg.		arc	*					efe	
<i>Vicia sepium</i>		nat	*					dvn	
<i>Vicia sylvatica</i>		nat	V				x	dvo	
<i>Vicia tenuifolia</i>		nat	3	x				bjk	
<i>Vicia tetrasperma</i>		nat	*					dvp	
<i>Vinca minor</i>		neo	*				x	dvs	
<i>Vincetoxicum hirundinaria</i>		nat	*	x				bjl	
<i>Viola arvensis</i>		arc	*					dvt	
<i>Viola canina</i>		nat	V	x				dvu	
<i>Viola hirta</i>		nat	*	x				dvv	
<i>Viola mirabilis</i>		nat	3				x	bjn	
<i>Viola odorata</i>		neo	*		x		x	dwv	
<i>Viola palustris</i>		nat	V	x				dvx	
<i>Viola persicifolia</i>		nat	2	x				bjo	
<i>Viola reichenbachiana</i>		nat	*				x	dvy	
<i>Viola riviniana</i>		nat	*				x	dvz	
<i>Viola tricolor</i>		nat	*	x				dwa	bjp
<i>Viola x bavaria</i>		nat	*				x	dwe	
<i>Viscum album</i>		nat	*					dwb	
<i>Vulpia bromoides</i>		nat	2	x				bjq	
<i>Vulpia myuros</i>		nat	*	x				dwc	
<i>Wolffia arrhiza</i>		nat	3					bjs	
<i>Woodsia ilvensis</i>		nat	1	x				bjt	
<i>x Calamophila baltica</i>	x	nat	*	x				dwk	
<i>x Festulolium loliaceum</i>		nat	*					faw	
<i>Xanthium albinum</i>		neo	*		x			dwd	
<i>Zannichellia palustris</i>		nat	V		x			bju	dwg, dwh
<i>Zostera marina</i>	x	nat	3					dwi	
<i>Zostera noltii</i>		nat	3					dwj	

Table A.3: Codes and full names of (sub-)species assigned to the species and/or aggregate of higher taxonomical level. Added to = Code of species to which the particular (sub)species was assigned to (see Table A.2 for full names of species).

Code	Species	Added to
116	<i>Taraxacum commixtum</i>	bgi
122	<i>Taraxacum discretum</i>	bgi
126	<i>Taraxacum franconicum</i>	bgi
14	<i>Sambucus nigra f. laciniata</i>	dln
144	<i>Polygonum arenastrum</i>	dag
181	<i>Taraxacum prionum</i>	eex
182	<i>Taraxacum rubrisquameum</i>	eex
191	<i>Nasturtium x sterile</i>	edb
206	<i>Asplenium trichomanes ssp. pachyrachis</i>	acw
224	<i>Asplenium trichomanes ssp. hastatum</i>	acw
225	<i>Festuca rubra ssp. juncea</i>	397
232	<i>Aster x salignus</i>	ear
238	<i>Luzula luzuloides ssp. rubella</i>	aub
240	<i>Festuca valesiaca ssp. parviflora</i>	ecc
243	<i>Epipactis helleborine ssp. neerlandica</i>	370
26	<i>Taraxacum gelertii</i>	eex
260	<i>Trichophorum cespitosum nothosp. foersteri</i>	bhg
27	<i>Taraxacum subalpinum</i>	bjg
276	<i>Polypodium x mantoniae</i>	edm
283	<i>Taraxacum leptoglotte</i>	eex
300	<i>Barbarea vulgaris ssp. arcuata</i>	ces
301	<i>Barbarea vulgaris ssp. rivularis</i>	ces
34	<i>Taraxacum rubicundum</i>	bgi
40	<i>Taraxacum paucilobum</i>	bjg
401	<i>Erigeron annuus ssp. strigosus</i>	cmx
413	<i>Festuca heteromalla</i>	397

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Table A.3 – Continued from previous page

Code	Species	Added to
50	<i>Aster novae-angliae</i>	ear
54	<i>Festuca nigrescens</i>	397
56	<i>Taraxacum disseminatum</i>	bgj
59	<i>Taraxacum bavaricum</i>	bgj
ace	<i>Armeria maritima</i> ssp. <i>elongata</i>	393
acf	<i>Armeria maritima</i> ssp. <i>halleri</i>	393
adt	<i>Bolboschoenus maritimus</i>	398
aes	<i>Callitriche cophocarpa</i>	eay
aeu	<i>Callitriche palustris</i> +	eay
agy	<i>Carex viridula</i> var. <i>pulchella</i>	agm
ahx	<i>Cerastium pumilum</i>	ebh
anj	<i>Festuca pallens</i>	ecc
ank	<i>Festuca valesiaca</i> ssp. <i>valesiaca</i>	ecc
aoj	<i>Galium verum</i> s.l.	cph
apr	<i>Helianthemum nummularium</i> ssp. <i>nummularium</i>	apq
aps	<i>Helianthemum nummularium</i> ssp. <i>obscurum</i>	apq
asf	<i>Lamium purpureum</i> var. <i>incisum</i>	ect
ate	<i>Lilium bulbiferum</i> ssp. <i>bulbiferum</i>	atd
atf	<i>Lilium bulbiferum</i> ssp. <i>croceum</i>	atd
avi	<i>Montia fontana</i> ssp. <i>chondrosperma</i>	avh
awm	<i>Ononis repens</i> ssp. <i>procurrens</i>	awn
awo	<i>Ononis spinosa</i>	awn
ayt	<i>Plantago major</i> ssp. <i>winteri</i>	edi
bbe	<i>Ranunculus aquatilis</i> +	bbd
bbg	<i>Ranunculus peltatus</i> ssp. <i>baudotii</i>	bbd
bbi	<i>Ranunculus circinatus</i>	bbd
bbj	<i>Ranunculus fluitans</i>	bbd
bbk	<i>Ranunculus hederaceus</i>	bbd
bbm	<i>Ranunculus nemorosus</i>	bbs
bbn	<i>Ranunculus oleucos</i>	bbd
bbo	<i>Ranunculus peltatus</i> ssp. <i>peltatus</i>	bbd
bbq	<i>Ranunculus polyanthemoides</i>	bbs
bbr	<i>Ranunculus polyanthemophyllus</i>	bbs
bbw	<i>Ranunculus serpens</i>	bbs
bbx	<i>Ranunculus trichophyllus</i>	bbd
bcq	<i>Rosa vosagiana</i> agg.	dds
bda	<i>Salicornia europaea</i> ssp. <i>brachystachya</i>	eee
bdd	<i>Salix repens</i> ssp. <i>repens</i>	eej
bdj	<i>Sanguisorba minor</i> ssp. <i>minor</i>	eel
bfk	<i>Sonchus arvensis</i> ssp. <i>uliginosus</i>	eer
bhh	<i>Trichophorum cespitosum</i> ssp. <i>cespitosum</i>	bhg
bhi	<i>Trichophorum cespitosum</i> ssp. <i>germanicum</i>	bhg
bif	<i>Utricularia vulgaris</i> +	bie
bim	<i>Valeriana sambucifolia</i>	efc
bin	<i>Valeriana wallrothii</i>	efc
bjp	<i>Viola tricolor</i> ssp. <i>curtisii</i>	dwa
cag	<i>Achillea millefolium</i>	eab
can	<i>Aethusa cynapium</i> ssp. <i>elata</i>	aec
cas	<i>Agrostis gigantea</i>	ead
cay	<i>Alchemilla filicaulis</i>	aaq
caz	<i>Alchemilla glabra</i>	aaq
cba	<i>Alchemilla micans</i>	aaq
cbb	<i>Alchemilla monticola</i>	aaq
cbe	<i>Alchemilla propinqua</i>	aaq
cbd	<i>Alchemilla subcrenata</i>	aaq
cbe	<i>Alchemilla subglobosa</i>	aaq
cbf	<i>Alchemilla vulgaris</i> +	aaq
cbg	<i>Alchemilla xanthochlora</i>	aaq
cbh	<i>Alisma plantago-aquatica</i>	ea
cbv	<i>Amaranthus blitum</i> ssp. <i>blitum</i>	363
cby	<i>Amaranthus blitum</i> ssp. <i>emarginatus</i>	363
cbz	<i>Amaranthus powellii</i>	eag
ccx	<i>Anthyllis vulneraria</i> ssp. <i>maritima</i>	391
ccy	<i>Anthyllis vulneraria</i> ssp. <i>pseudovulneraria</i>	391
cdl	<i>Arenaria serpyllifolia</i> ssp. <i>serpyllifolia</i>	ean
cdm	<i>Armeria maritima</i> ssp. <i>maritima</i>	393
cdw	<i>Aster lanceolatus</i>	ear
cdx	<i>Aster novi-belgii</i> +	ear
cdy	<i>Aster parviflorus</i>	ear
cei	<i>Atriplex prostrata</i>	eas
cep	<i>Ballota nigra</i> ssp. <i>meridionalis</i>	adj
ceq	<i>Ballota nigra</i> ssp. <i>nigra</i>	adj
cey	<i>Betula pubescens</i> ssp. <i>carpatica</i>	eat
cez	<i>Betula pubescens</i> ssp. <i>pubescens</i>	eat
cfm	<i>Bromus hordeaceus</i>	eau
cfz	<i>Callitriche hamulata</i>	eay

Continued on next page

Table A.3 – Continued from previous page

Code	Species	Added to
cga	<i>Callitriche obtusangula</i>	eay
cgb	<i>Callitriche platycarpa</i>	eay
cgc	<i>Callitriche stagnalis</i>	eay
cgg	<i>Cardamine pratensis</i>	eaz
cgy	<i>Carex arenaria</i>	ebc
chf	<i>Carex guestphalica</i>	ebe
chk	<i>Carex pairae</i>	ebe
chs	<i>Carex spicata</i>	ebe
chw	<i>Carex vulpina</i>	ebg
cih	<i>Centaurium littorale</i> ssp. <i>littorale</i>	ahn
cii	<i>Centaurium littorale</i> ssp. <i>uliginosum</i>	ahn
cjc	<i>Chenopodium rubrum</i>	ebj
cjd	<i>Chenopodium strictum</i>	ebi
cje	<i>Chenopodium sueticum</i>	ebi
cka	<i>Crataegus laevigata</i> ssp. <i>laevigata</i>	ebl
ckb	<i>Crataegus laevigata</i> ssp. <i>palmstruchii</i>	ebl
cke	<i>Crataegus monogyna</i> ssp. <i>monogyna</i>	ebo
ckf	<i>Crataegus monogyna</i> ssp. <i>nordica</i>	ebo
cmh	<i>Epilobium tetragonum</i> ssp. <i>lamyi</i>	ebw
cmj	<i>Epipactis helleborine</i> ssp. <i>helleborine</i>	370
cmk	<i>Epipactis leptochila</i> ssp. <i>leptochila</i>	amj
cml	<i>Epipactis leptochila</i> ssp. <i>neglecta</i>	amj
cnq	<i>Euphrasia micrantha</i>	anf
cnr	<i>Euphrasia nemorosa</i> +	anf
cnx	<i>Festuca filiformis</i>	ecc
cnz	<i>Festuca guestfalica</i>	ecc
coa	<i>Festuca ovina</i> +	ecc
cod	<i>Festuca rubra</i> ssp. <i>arenaria</i>	397
coe	<i>Festuca rubra</i> ssp. <i>litoralis</i>	397
cog	<i>Festuca rupicola</i>	ecc
coh	<i>Festuca brevipila</i>	ecc
coi	<i>Filipendula ulmaria</i> var. <i>denudata</i>	ece
coj	<i>Filipendula ulmaria</i> var. <i>ulmaria</i>	ece
coq	<i>Fumaria officinalis</i> ssp. <i>wirtgenii</i>	ecg
csz	<i>Lamium galeobdolon</i>	ecq
cta	<i>Lamium montanum</i>	ecq
ctr	<i>Leonurus cardiaca</i> ssp. <i>cardiaca</i>	asy
cts	<i>Leonurus cardiaca</i> ssp. <i>villosus</i>	asy
ctx	<i>Galium palustre</i> ssp. <i>elongatum</i>	cpc
cub	<i>Leucanthemum ircutianum</i>	ecu
cwp	<i>Monotropa hypophegea</i>	avg
cwq	<i>Monotropa hypopitys</i> +	avg
cwr	<i>Montia fontana</i> ssp. <i>amporitana</i>	avh
cws	<i>Montia fontana</i> ssp. <i>fontana</i>	avh
cwt	<i>Montia fontana</i> ssp. <i>variabilis</i>	avh
cxg	<i>Nasturtium microphyllum</i>	edb
cyi	<i>Papaver dubium</i> ssp. <i>dubium</i>	ede
cyj	<i>Papaver dubium</i> ssp. <i>lecoqii</i>	ede
cyk	<i>Polygala vulgaris</i> ssp. <i>collina</i>	aze
cyq	<i>Polygala vulgaris</i> ssp. <i>oxyptera</i>	aze
cza	<i>Phyteuma spicatum</i> ssp. <i>coeruleum</i>	ayo
czb	<i>Phyteuma spicatum</i> ssp. <i>spicatum</i>	ayo
czk	<i>Plantago major</i> ssp. <i>intermedia</i>	edi
czl	<i>Plantago major</i> ssp. <i>major</i>	edi
daa	<i>Polygala amara</i> ssp. <i>brachyptera</i>	azc
dab	<i>Polygala amarella</i>	azc
dal	<i>Persicaria lapathifolia</i> ssp. <i>brittingeri</i>	edl
dam	<i>Persicaria lapathifolia</i> ssp. <i>pallida</i>	edl
dar	<i>Polypodium interjectum</i>	edm
daw	<i>Potamogeton berchtoldii</i>	edo
dco	<i>Ranunculus penicillatus</i>	bbd
ddf	<i>Rosa caesia</i>	bcl
ddm	<i>Rosa pseudoscabriuscula</i>	bcp
ddn	<i>Rosa sherardii</i>	bcp
ddq	<i>Rosa tomentosa</i> +	bcp
ddr	<i>Rosa villosa</i>	bcp
dkh	<i>Sagina micropetala</i>	eed
dkn	<i>Salicornia europaea</i> ssp. <i>europaea</i>	eee
dkq	<i>Salix alba</i> ssp. <i>vitellina</i>	eeg
dkv	<i>Salix fragilis</i> +	eeh
dky	<i>Salix purpurea</i> ssp. <i>lambertiana</i>	eei
dla	<i>Salix repens</i> ssp. <i>dunensis</i>	eej
dlb	<i>Salix triandra</i> ssp. <i>amygdalina</i>	eek
dle	<i>Salix x rubens</i>	eeh
dlg	<i>Salsola kali</i> ssp. <i>kali</i>	bdf
dlh	<i>Salsola kali</i> ssp. <i>tragus</i>	bdf

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Table A.3 – Continued from previous page

Code	Species	Added to
dlw	<i>Scleranthus polycarpus</i>	een
dmd	<i>Sedum maximum</i>	eeo
dme	<i>Sedum telephium</i> +	eeo
dnm	<i>Solanum nigrum</i> ssp. <i>schultesii</i>	eeq
dnw	<i>Sonchus arvensis</i> ssp. <i>arvensis</i>	eer
doc	<i>Sparganium erectum</i> ssp. <i>neglectum</i>	ees
dpo	<i>Taraxacum balticum</i>	bgj
dpq	<i>Taraxacum brachyglossum</i>	bgj
dpr	<i>Taraxacum bracteatum</i>	eex
dpx	<i>Bolboschoenus maritimus</i> x <i>yagara</i>	398
dqj	<i>Taraxacum haematicum</i>	eex
dqv	<i>Taraxacum lacistophyllum</i>	bgj
drq	<i>Taraxacum nordstedtii</i>	eex
drn	<i>Taraxacum palustre</i>	bgj
drw	<i>Taraxacum proximum</i>	bgj
dsa	<i>Taraxacum duplidentifrons</i>	eex
dsd	<i>Taraxacum scanicum</i>	bgj
dsg	<i>Taraxacum parnassicum</i>	bgj
dsr	<i>Taraxacum tenuilobum</i>	bgj
dss	<i>Taraxacum tortilobum</i>	bgj
dsx	<i>Festuca polesica</i>	ecc
duu	<i>Veronica catenata</i>	biv
duy	<i>Veronica hederifolia</i> ssp. <i>lucorum</i>	efd
dwg	<i>Zannichellia palustris</i> ssp. <i>palustris</i>	bju
dwh	<i>Zannichellia palustris</i> ssp. <i>pedicellata</i>	bju
eah	<i>Anagallis arvensis</i> var. <i>caerulea</i>	ckc
eap	<i>Asplenium trichomanes</i> ssp. <i>trichomanes</i>	acw
ebk	<i>Crataegus curvisepala</i> agg.	cjz
ebm	<i>Crataegus laevigata</i>	eb1
ebq	<i>Dactylorhiza fuchsii</i>	akr
ebu	<i>Eleocharis palustris</i> ssp. <i>palustris</i>	clr
ebv	<i>Eleocharis palustris</i> ssp. <i>vulgaris</i>	clr
ecc	<i>Festuca rubra</i>	397
ecm	<i>Hypericum maculatum</i> ssp. <i>obtusiusculum</i>	crp

Table A.4: Soil types listed in the soil map of Lower Saxony (BÜK 50) and the assignment to 27 soil type categories. Abbr. = Abbreviation of soil type.

Abbr.	Soil type	Category
AB	Vega	1
AB-S	Vega-Pseudogley	1
B-P	Braunerde-Podsol	2
B	Braunerde	2
B-Q	Braunerde-Regosol	2
B-D	Braunerde-Pelosol	2
B-S	Braunerde-Pseudogley	2
B-R	Braunerde-Rendzina	2
B-pL	Braunerde-Parabraunerde; podsoliert	2
B-N	Braunerde-Ranker	2
B-Z	Braunerde-Pararendzina	2
B-pQ	Braunerde-Regosol; podsoliert	2
B-L	Braunerde-Parabraunerde	2
D	Pelosol	3
D-S	Pelosol-Pseudogley	3
D-B	Pelosol-Braunerde	3
E//P-G	Plaggenesch unterlagert von Podsol-Gley	4
E//P	Plaggenesch unterlagert von Podsol	4
E/G-P	Gley-Podsol mit Plaggenauflage	4

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Table A.4 – Continued from previous page

Abbr.	Soil type	Category
E//G	Plaggenesch unterlagert von Gley	4
E/S-P	Pseudogley-Podsol mit Plaggenauflage	4
E//S	Plaggenesch unterlagert von Pseudogley	4
E//S-B	Plaggenesch unterlagert von Pseudogley-Braunerde	4
E//B	Plaggenesch unterlagert von Braunerde	4
E/P-B	Podsol-Braunerde mit Plaggenauflage	4
E	Plaggenesch	4
E/S-B	Pseudogley-Braunerde mit Plaggenauflage	4
E/B	Braunerde mit Plaggenauflage	4
E//G-B	Plaggenesch unterlagert von Gley-Braunerde	4
E/G-B	Gley-Braunerde mit Plaggenauflage	4
E//B-G	Plaggenesch unterlagert von Braunerde-Gley	4
E//S-L	Plaggenesch unterlagert von Pseudogley-Parabraunerde	4
E/L	Parabraunerde mit Plaggenauflage	4
E//L	Plaggenesch unterlagert von Parabraunerde	4
E/S	Pseudogley mit Plaggenauflage	4
E/P	Podsol mit Plaggenauflage	4
E//P-B	Plaggenesch unterlagert von Podsol-Braunerde	4
E//S-P	Plaggenesch unterlagert von Pseudogley-Podsol	4
G	Gley	5
G-P	Gley-Podsol	5
G-S	Gley-Pseudogley	5
G-Q	Gley-Regosol	5
G-AB	Gley-Vega	5
G-B	Gley-Braunerde	5
G-T	Gley-Schwarzerde	5
G-OL	Gley-Lockersyrosem	5
G-P-S	Gley-Podsol-Pseudogley	5
G-L	Gley-Parabraunerde	5
G-Z	Gley-Pararendzina	5
HHv	Erd-Hochmoor	6
HNv/G	Gley mit Erd-Niedermoorauflage	6
HNv	Erd-Niedermoor	6
HHv/G-P	Gley-Podsol mit Erd-Hochmoorauflage	6
HH	Hochmoor	6
HHv//S-G	Erd-Hochmoor unterlagert von Pseudogley-Gley	6
HHv/G	Gley mit Erd-Hochmoorauflage	6
K//G	Kolluvisol unterlagert von Gley	7
K//L	Kolluvisol unterlagert von Parabraunerde	7
K//S-L	Kolluvisol unterlagert von Pseudogley-Parabraunerde	7
K//B	Kolluvisol unterlagert von Braunerde	7
K	Kolluvisol	7
L	Parabraunerde	8
L-S	Parabraunerde-Pseudogley	8
L-SH	Parabraunerde-Haftäössepseudogley	8
pL	Parabraunerde; podsoliert	8
MC	Kalkmarsch	9
MD	Dwogmarsch	10

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Table A.4 – *Continued from previous page*

Abbr.	Soil type	Category
MK//HN	Knickmarsch unterlagert von Niedermoor	11
MK	Knickmarsch	11
MK/G	Gley mit Knickaufflage	11
MK//HH	Knickmarsch unterlagert von Hochmoor	11
MN	Kleimarsch	12
MN//HN	Kleimarsch unterlagert von Niedermoor	12
MN/HN	Niedermoor mit Knickmarschaufflage	12
MN//HH	Kleimarsch unterlagert von Hochmoor	12
MN/HH	Hochmoor mit Knickmarschaufflage	12
MO	Organomarsch	13
MO//HN	Organomarsch unterlagert von Niedermoor	13
MR	Rohmarsch	14
MR/HN	Niedermoor mit Organomarschaufflage	14
MR/HH	Hochmoor mit Rohmarschaufflage	14
N	Ranker	15
O	Syrosem	16
OL	Lockersyrosem	16
P-G	Podsol-Gley	17
P	Podsol	17
P-B	Podsol-Braunerde	17
P-S	Podsol-Pseudogley	17
P-Q	Podsol-Regosol	17
Q	Regosol	18
pQ	podsoliger Regosol	18
S-P	Pseudogley-Podsol	19
S-B	Pseudogley-Braunerde	19
S	Pseudogley	19
S-G	Pseudogley-Gley	19
S-L	Pseudogley-Parabraunerde	19
S-T	Pseudogley-Schwarzerde	19
S//MN	Pseudogley unterlagert von Kleimarsch	19
S-AB	Pseudogley-Vega	19
S-D	Pseudogley-Pelosol	19
S-K	Pseudogley-Kolluvisol	19
pS	Pseudogley; podsoliert	19
SH	Haftnössepseudogley	19
S-Q	Pseudogley-Regosol	19
R	Rendzina	20
F	Felshumusboden	21
T-L	Schwarzerde-Parabraunerde	22
T	Schwarzerde	22
YD\HH	Hochmoor mit Deckkulturbodenaufflage	23
YD/HH	Hochmoor mit Deckkulturbodenaufflage	23
YT	Spittmarsch	24
YUhn	Tiefumbruchboden	25
YUp	Tiefumbruchboden	25
YUg	Tiefumbruchboden	25
YUhh	Tiefumbruchboden	25

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Table A.4 – Continued from previous page

Abbr.	Soil type	Category
YUg-p	Tiefumbruchboden	25
YUh	Tiefumbruchboden	25
YUs-p	Tiefumbruchboden	25
YUmn	Tiefumbruchboden	25
YUab	Tiefumbruchboden	25
YUs-ab	Tiefumbruchboden	25
YUg-ab	Tiefumbruchboden	25
YUs	Tiefumbruchboden	25
YUs-g	Tiefumbruchboden	25
Z	Pararendzina	26
ZS	Strand	27

Table A.5: Roads that are represented as polylines in the ATKIS-DLM were buffered by using the buffer distance specified below. Standard values for the width of roads was taken from FGSV (1996). n/a = not available.

Road category	FGSV (1996)	Buffer
Motorways (Autobahnen)	Minimum: 20 m Maximum: 29 m Standard: 23 m	11.5 m → $2 \times 11.5 \text{ m} = 23 \text{ m width}$
Federal highways (Bundesstraßen)	Minimum: 7.5 m Maximum: 11.5 m, 15 m for twin-track roads Standard: n/a.	5.625 m → $2 \times 5.625 \text{ m} = 11.25 \text{ m width}$ (average of 7.5 m and 15 m)
State roads (Landesstraßen)	Standard: 6.5 m	3.25 m → $2 \times 3.25 \text{ m} = 6.5 \text{ m width}$
County roads (Kreisstraßen)	n/a; standard of 6.5 m as- sumed	3.25 m → $2 \times 3.25 \text{ m} = 6.5 \text{ m width}$
Municipal roads (Gemeindestraßen)	n/a; standard of 5.5 m as- sumed	2.75 m → → $2 \times 2.75 \text{ m} = 5.5 \text{ m width}$

Table A.6: Land cover types (LCT) used for the calculation of landscape metrics, their description as well as the ATKIS-DLM objects included in the particular LCT.

LCT	Description	Objects ATKIS-DLM
Industrial areas	Areas used for (non agricultural) production (e.g. supply and disposal facilities, exhibition centers, mining areas etc.)	Abfallbeseitigungsanlage, Abfalldeponie, Absetzbecken/ Rieselfeld/ Schlammteich, Ausstellungsgelände/ Messegelände, Bergbaubetrieb, Fläche besonderer funktionaler Prägung, Förderanlage, Freileitung, Gärtnerei, Heizwerk, Industrie- und Gewerbefläche, Kläranlage/Klärwerk, Kraftwerk, Raffinerie, Tagebau/ Grube/ Steinbruch, Umspannwerk, Wasserwerk, Werft, Dock, Hafenbecken
Urban areas	Areas used for housing and recreation (e.g. residential areas, parks, campgrounds etc.)	Campingplatz, Denkmal/ Denkstein/ Standbild, Fläche gemischter Nutzung, Freilichtmuseum, Freilichttheater, Freizeitanlage, Freizeitpark/ Safari-park, Friedhof, Gartenland, Golfplatz, Grünanlage, Ortslage, Schiesstand, Schwimmbad/ Freibad, Schwimmbecken, Spielfeld/ Spielfläche, Sportanlage, Stadion, Wohnbaufläche, Zoo, Zuschauertribüne
Traffic areas	Areas used for transport (roads, railway lines, airports, etc.)	Bahnhofsanlage, Bahnkörper, Bahnstrecke, Brücke/ Unterführung/ Überführung, Fahrbahn, Flughafen, Flughafen-Vorfeld, Flugplatz/ Landeplatz, Hafen, Hafenbecken, Platz (Rastplatz/ Parkplatz), Raststätte, Rollbahn, Schienenbahn, Schleuse, Schleusen-kammer, Straße, Straßenkörper, Tunnel, Weg
Croplands	Areas used for agricultural production	Ackerland, Rieselfeld, Sonderkultur
Forests	Forested areas	Wald/ Forst
Grasslands	Grassland areas	Grünland
Water bodies	Standing and flowing waters (e.g. lakes, streams, channels, springs)	Binnensee/ Stausee/ Teich, Graben/ Kanal (Wasserwirtschaft), Kanal (Schifffahrt), Strom/ Fluss/ Bach, Talsperre/ Wehr, Quelle
Others	Areas not belonging to any of the above mentioned LCT (e.g. tree rows, bogs, islands)	Fläche z.Zt. unbestimmbar, Grenze, Halde, Aufschüttung, Nasser Boden, Vegetationslose Fläche, Zaun, Uferbefestigung, Heide, Baumgruppe/ Baumreihe, Gehölz, Moor, Insel, Meer, Priel, Sandbank, Watt

Table A.7: Characterisation of the landscapes in the six clusters (mean value of the standardised variable per cluster \pm SE) found for 1383 landscape units in Lower Saxony, Germany. *n*, number of landscape units in the particular cluster. Superscript letters indicate significant differences between clusters ($p \leq 0.05$, non-parametric Kruskal-Wallis test).

Variable names	A	B	C	D	E	F
<i>n</i>	131	383	148	389	242	90
Landscape composition						
prop_water	0.84 \pm 0.10 ^a	0.02 \pm 0.04 ^b	1.36 \pm 0.10 ^a	-0.57 \pm 0.02 ^c	-0.18 \pm 0.05 ^d	-0.58 \pm 0.07 ^c
prop_forest	-0.53 \pm 0.08 ^a	0.14 \pm 0.03 ^b	-1.73 \pm 0.05 ^c	0.32 \pm 0.04 ^d	0.08 \pm 0.05 ^b	1.44 \pm 0.02 ^e
prop_grass	-0.12 \pm 0.09 ^a	0.16 \pm 0.03 ^b	1.46 \pm 0.04 ^c	-0.70 \pm 0.03 ^d	0.57 \pm 0.04 ^e	-1.38 \pm 0.07 ^f
prop_crop	-0.58 \pm 0.10 ^a	0.35 \pm 0.02 ^b	-0.80 \pm 0.08 ^{ac}	0.59 \pm 0.02 ^d	-0.01 \pm 0.03 ^e	-1.87 \pm 0.16 ^c
prop_urban	1.69 \pm 0.07 ^a	0.44 \pm 0.03 ^b	0.02 \pm 0.05 ^c	-0.27 \pm 0.03 ^d	-0.58 \pm 0.04 ^e	-1.65 \pm 0.09 ^f
prop_industry	1.75 \pm 0.06 ^a	0.19 \pm 0.04 ^b	-0.44 \pm 0.06 ^c	-0.43 \pm 0.03 ^c	-0.02 \pm 0.07 ^d	-0.75 \pm 0.07 ^e
prop_traffic	1.92 \pm 0.10 ^a	0.36 \pm 0.03 ^b	-0.06 \pm 0.07 ^c	-0.31 \pm 0.03 ^d	-0.61 \pm 0.03 ^c	-1.24 \pm 0.05 ^f
prop_others	0.60 \pm 0.06 ^a	-0.27 \pm 0.03 ^b	-0.29 \pm 0.06 ^b	-0.62 \pm 0.02 ^c	1.24 \pm 0.06 ^d	0.07 \pm 0.16 ^b
all_SHDI	1.19 \pm 0.05 ^a	0.50 \pm 0.03 ^b	-0.73 \pm 0.06 ^c	-0.55 \pm 0.03 ^c	0.54 \pm 0.03 ^b	-1.76 \pm 0.11 ^d
Landscape configuration						
meshsize	-1.35 \pm 0.02 ^a	-0.48 \pm 0.03 ^b	-0.23 \pm 0.07 ^b	0.40 \pm 0.04 ^c	0.50 \pm 0.05 ^c	1.31 \pm 0.08 ^d
all_Area_Am	-0.81 \pm 0.06 ^a	-0.70 \pm 0.03 ^a	0.31 \pm 0.07 ^b	0.61 \pm 0.03 ^c	-0.29 \pm 0.04 ^d	1.79 \pm 0.07 ^e
all_Shape_Am	0.93 \pm 0.12 ^a	-0.25 \pm 0.03 ^b	1.39 \pm 0.11 ^a	-0.24 \pm 0.03 ^{bc}	-0.38 \pm 0.03 ^{bc}	-0.48 \pm 0.06 ^c
for_NP	-0.35 \pm 0.07 ^a	0.69 \pm 0.05 ^b	-0.99 \pm 0.04 ^c	-0.12 \pm 0.04 ^a	0.21 \pm 0.05 ^d	-0.87 \pm 0.04 ^c
grass_NP	0.24 \pm 0.08 ^a	0.82 \pm 0.05 ^b	-0.61 \pm 0.07 ^c	-0.21 \pm 0.04 ^d	-0.24 \pm 0.04 ^d	-1.27 \pm 0.06 ^e
crop_NP	-0.07 \pm 0.08 ^a	0.68 \pm 0.05 ^b	0.48 \pm 0.10 ^c	-0.60 \pm 0.02 ^d	0.12 \pm 0.04 ^{ac}	-1.32 \pm 0.05 ^e

Table A.8: Overlap of species sub-groups for which the richness patterns along the gradient of landscape modification intensity for 1383 landscape units in Lower Saxony, Germany, were analysed. Total species numbers for the particular sub-groups are printed in bold. In total, 1594 species were considered in the analysis.

	Native species	Established neophytes	Threatened species	Forest species	Urban species	EIV species N1-3	EIV species N7-9
Native species	1297						
Established neophytes	–	143					
Threatened species	483	9	538				
Forest species	188	9	65	197			
Urban species	12	32	10	1	68		
EIV species N1-3	444	16	286	36	6	474	
EIV species N7-9	218	47	59	56	33	–	306

Table A.9: Species richness per cluster for the total number of species occurring in 1383 landscape units in Lower Saxony, Germany, as well as seven sub-groups according to their residence status, threat status and habitat preferences. Superscript letters indicate significant differences between clusters ($p \leq 0.05$, non-parametric Kruskal-Wallis test).

	A	B	C	D	E	F
All species	528.0±8.3 ^a	461.3±4.1 ^b	362.7±5.6 ^c	460.5±4.4 ^b	414.7±4.8 ^d	442.8±8.4 ^{bcd}
Native species	399.9±6.1 ^a	365.6±3.3 ^b	285.2±4.6 ^c	364.8±3.6 ^b	335.2±3.7 ^d	365.0±7.2 ^b
Established neophytes	50.7±1.2 ^a	35.7±0.4 ^b	29.2±0.7 ^c	32.0±0.4 ^d	28.9±0.5 ^c	27.1±0.9 ^c
Threatened species	32.8±1.5 ^a	27.2±0.8 ^{bc}	17.9±1.0 ^d	26.8±0.9 ^b	29.6±1.0 ^{ac}	34.3±1.9 ^a
Forest species	53.9±1.9 ^{ab}	50.3±1.1 ^a	22.9±1.0 ^c	59.9±1.4 ^b	40.9±1.0 ^d	59.5±3.3 ^{ab}
Urban species	32.1±0.8 ^a	22.5±0.3 ^b	17.9±0.4 ^c	22.7±0.3 ^b	17.5±0.4 ^c	16.9±0.6 ^c
EIV species N1-3	98.2±2.3 ^{ab}	95.3±1.2 ^a	57.8±2.1 ^c	88.5±1.2 ^d	96.0±1.4 ^a	108.3±2.3 ^b
EIV species N7-9	148.6±2.3 ^a	125±1.1 ^b	113±1.4 ^c	126.2±1.2 ^b	106.6±1.2 ^c	107.0±2.7 ^c

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Appendix B

The contents of Appendix B can be found on the DVD accompanying this thesis.

PDF file of the present dissertation

Diss_Schmiedel.pdf

Schmiedel, I. (2014): Vascular plant species richness at the landscape scale: Patterns and processes. Dissertation. Georg-August-University of Göttingen. 218 pp.

PDF files of publications related to this thesis

Schmidt_et_al_2014.pdf

Schmidt, M.; Mölder, A.; Schönfelder, E.; Engel, F.; Schmiedel, I. & Culmsee, H. (2014): Determining ancient woodland indicator plants for practical use: A new approach developed in northwest Germany. *Forest Ecology and Management* **330**: 228–239.

Culmsee_et_al_2014.pdf

Culmsee, H.; Schmidt, M.; Schmiedel, I.; Schacherer, A.; Meyer, P. & Leuschner, C. (2014): Predicting the distribution of forest habitat types using indicator species to facilitate systematic conservation planning. *Ecological Indicators* **37**: 131–144.

Schmiedel_et_al_2013.pdf

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Schmidt_et_al_2011.pdf

Schmidt, M.; Culmsee, H.; Boch, S.; Heinken, T.; Müller, J. & Schmiedel, I. (2011): Anwendungsmöglichkeiten von Waldartenlisten für Gefäßpflanzen, Moose und Flechten. *BfN-Skripten* 299: 25–45.

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Schmiedel, I.; Schacherer, A.; Hauck, M.; Schmidt, M. & Culmsee, H. (2011): Verbreitungsmuster der Farn- und Blütenpflanzen in Niedersachsen und Bremen unter Berücksichtigung ihres Einbürgerungsstatus und ihrer Gefährdungssituation. *Tuexenia* 31: 211–226.