Rewetting does not return drained fen peatlands to their old selves

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Peatlands have been drained for land use for a long time and on a large scale, turning them from carbon and nutrient sinks into respective sources, diminishing water regulation capacity, causing surface height loss and destroying biodiversity. Over the last decades, drained peatlands have been rewetted for biodiversity restoration and, as it strongly decreases greenhouse gas emissions, also for climate protection. We quantify restoration success by comparing 320 rewetted fen peatland sites to 243 near-natural peatland sites of similar origin across temperate Europe, all set into perspective by 10k additional European fen vegetation plots. Results imply that rewetting of drained fen peatlands induces the establishment of tall, graminoid wetland plants (helophytisation) and long-lasting differences to pre-drainage biodiversity (vegetation), ecosystem functioning (geochemistry, hydrology), and land cover characteristics (spectral temporal metrics). The Paris Agreement entails the rewetting of 500,000 km² of drained peatlands worldwide until 2050-2070. A better understanding of the resulting locally novel ecosystems is required to improve planning and implementation of peatland rewetting and subsequent management.
already in 1658, the world’s first scientific book on peatlands included a chapter on peatland restoration and addressed the question of their long-term use. However, centuries of peatland destruction followed and large-scale restoration of drained peatlands is just about to begin. Despite some progress in understanding ecological functioning of rewetted peatlands, the early warning from 1764 that restoration success may be slow can only now be quantified at a substantial amount of rewetted sites, across continental scales and multiple ecosystem functions.

Intact peatlands provide numerous ecosystem services. They store huge amounts of carbon (600 ± 100 Gt), thus 30% of the global soil carbon is found on only 3% of the global land, they regulate water quality and quantity, and they harbor highly specialized biota. Approx. 500,000 km², i.e., 10–15% of the current peatland area, have been drained for agriculture, peat extraction and forestry, historically mainly in temperate and boreal regions, but more recently also in the tropics. Drainage of peatlandsimpairs their ecosystem service provisioning. Peat oxidation in drained peatlands is responsible for about 5% of the global anthropogenic greenhouse gas (GHG) emissions. Peat mineralization leads to ongoing land subsidence and turns drained peatlands into sources of nutrients, causing eutrophication of downstream surface and groundwater. A fundamental distinction is commonly made between bogs that are solely fed by rainwater, and fens that also receive ground- and/or surface water that has been in contact with the mineral soil or bedrock. Fens cover more than half of Europe’s peatland area. Especially temperate fens, which often become nutrient-rich and productive upon drainage, have been drained for agriculture. Consequently, many temperate fen peatland species are globally endangered because of habitat loss.

Rewetting drained peatlands can strongly reduce or stop net carbon loss immediately and may lead to new carbon sequestration. In contrast, continued emissions from drained peatlands may comprise 12–41% of the GHG emission budget. Peat oxidation during drainage has altered peat physical parameters and has led to, for instance, increased bulk density and decreased porosity, hydraulic conductivity and storativity, leading to stronger water table fluctuations. Periodic or episodic inundation can provoke peaks of methane emissions. Furthermore, nutrient availability after rewetting is considerably higher than in natural peatlands, mainly because of peat mineralization and fertilization while the peatland was drained, and because of mobilization of phosphorus upon rewetting. Consequently, microbial and plant communities show prompt recovery towards their pre-drainage status only in previously weakly disturbed ecosystems.

Here, we compare 320 rewetted fen sites with 243 near-natural sites of similar origin from the major fen peatland regions of Europe (Fig. 1), which are further set into perspective by additional 10k vegetation plots, randomly chosen from >90k plots of European fen vegetation and land cover characteristics. Our data imply that rewetting drained fens induces the establishment of tall, graminoid wetland plants, i.e., a helophytisation, with no trend back to their former biodiversity and ecosystem functioning for at least several decades in more than half of the sites. A better understanding of biodiversity and ecological functions of these locally novel ecosystems is urgently required to assess restoration success and to improve planning and implementation of peatland rewetting and the subsequent management.

Results and discussion
Rewetted peatlands differ in their biodiversity and ecosystem functioning from near-natural peatlands. Rewetted fen peatlands differ in biodiversity and ecosystem functioning from near-natural peatlands, in particular regarding plant community composition and geochemistry (Fig. 2a, c). Smaller, but still highly significant differences were detected in hydrology and in land cover characteristics (Fig. 2b, d). The vegetation of the rewetted sites was furthermore less diverse at the plot scale across the range of Hill’s diversity numbers (Supplementary Fig. 1; Shannon diversity: 1.46 ± 0.04 (mean ± SE) in rewetted sites; 1.75 ± 0.04 in near-natural sites; p\textsubscript{\text{trend}} < 0.001).

The differences in plant community composition between rewetted and near-natural sites are frequently linked to a helophytisation, i.e., the shift to a dominance of tall, graminoid wetland plants, as indicated by significant preferences of tall helophytes such as Typha latifolia or Phalaris arundinacea for the rewetted sites (see Indicator Species Analysis in Supplementary Data 2), by a higher prevalence of the EUNIS 2020 habitat type “tall helophyte bed” (relative frequency in rewetted peatlands: 25.5% versus 6.2% in near-natural peatlands; Chi²: p < 0.001; Supplementary Table 1), and by a 66% higher cover sum of tall helophyte species in rewetted as compared to near-natural sites (Fig. 3). This helophytisation has implications beyond biodiversity as plant species composition affects carbon cycling by litter quality (e.g., polyphenol content), root exudates, and production and consumption of carbon in the rhizosphere, both of which impact gaseous emissions. The tall helophytes are highly conductive for gases due to their aerenchyma, which may increase or suppress methane emissions. Brown mosses with high peat formation potential are largely absent from rewetted peatlands dominated by helophytes (Fig. 3) and Supplementary Data 2). While rewetting clearly reduces carbon emissions by inhibiting peat mineralization, comparative analyses between the resulting vastly contrasting vegetation types are required to assess the total greenhouse gas effects of rewetting.

The helophytisation after rewetting is presumably caused by altered water table dynamics and geochemistry of rewetted peatlands. Peat mineralization, consolidation and compaction during the drained phase cause subsidence, leading to inundation after rewetting (mean median water table being 5.0 cm above surface in rewetted sites versus 1.5 cm below surface in near-natural sites and maximum water table differing by 74%; Fig. 3a, c). Peat mineralization is reflected in the lower organic matter content of the top soil layer (−18%) and in the higher bulk density (+61%; Fig. 3f, e), both of which decrease porosity and hydraulic conductivity, provoking larger water table amplitudes (+15%) in rewetted sites (Fig. 3b). To sum up, inundation and eutrophic conditions promote tall helophytes, and these suppress low vascular plants and mosses by light competition.

Land cover characteristics, i.e., the 208 metrics of the spectral-temporal variability of land surfaces in Earth observation time series, also show differences between rewetted and near-natural peatlands (Fig. 2d), mainly attributed to high variety in the modified normalized difference water index (MNDWI) and the near infrared reflectance (nIR) for rewetted surfaces (Supplementary Table 2). High MNDWI values relate to a high share of open water in the pixels, nIR values correspond to vegetation.
density, vigor or type and are lowest for open water. Thus, both metrics relate to differences of the water regime for the vegetated surfaces. The dissimilarity between land cover characteristics in rewetted sites and near-natural sites hints at different pathways after rewetting. Environmental conditions such as hydrology, (chance) priority effects of colonizing species or former management - the rewetted sites did not differ from near-natural sites in share of unused sites after rewetting (Fig. 3l) - may determine the outcome.

A basic feature of natural peatlands is their stability, which is commonly attributed to a high degree of self-regulation, for instance with regards to buffering hydrological extremes by surface oscillation. Further, stable, azonal species compositions indicate some degree of independence from zonal climate conditions. Plant community composition and carbon dynamics in rewetted peatlands, however, are highly responsive to weather extremes, hinting at a reduced stability. In our dataset, rewetted peatlands in all four response clusters were generally more variant than near-natural peatlands (Fig. 2 insets). This variance of rewetted peatlands may have resulted from a poorly restored hydrology on the landscape level and potentially impairs their ecosystem functioning and service provisioning. For instance, reduced oscillation capacity creates both times with water tables dropping low and inducing carbon dioxide emissions due to increased mineralization, and times of inundation with increased methane emissions.

No general trend towards natural conditions up to three decades after rewetting. A strong and, apparently, long-lasting difference in biodiversity, ecosystem functioning, and land cover characteristics between each rewetted site and its closest near-natural counterpart exists (Fig. 4). Even though 40% (vegetation) to 80% (hydrology) of the rewetted sites generally resembled composition or functioning of typical near-natural peatlands (falling into the 95% confidence ellipses in Fig. 2), the rewetted sites differed strongly from near-natural counterparts in biodiversity, ecosystem functioning (hydrology, geochemistry) and land cover characteristics (linear models with $p_{\text{intercept}} < 0.001$ for all four response clusters despite low confidence due to small sample size for geochemistry). No detectable trend towards increasing similarity over time (linear models with $p_{\text{slope}} > 0.05$ for all four response clusters except vegetation with $p_{\text{slope}} = 0.013$ but still a predicted dissimilarity of 0.85 after 5 decades (95%CI: 0.64 to 1.07); Fig. 4) indicates that restoration success is either reached promptly after rewetting or not at all within the observed three decades covered by our data. Spatial variance exists also between near-natural counterparts of similar origin and distance, but at a much lower level than between rewetted and near-natural counterparts (horizontal dashed lines in Fig. 4 which is not included in the 95% confidence bands except for geochemistry with its low sample size). In short, time had little to no effect on dissimilarity between rewetted sites and their near-natural counterparts in our data, i.e., time seems to have little to no effect on restoration success for at least three decades.

It is commonly assumed that the stabilization of the water table close to the surface is a premise for successful restoration of peatlands because too low water tables lead to high carbon...
emissions and non-wetland plants remaining dominant while too high water tables create shallow lakes rather than peatlands. While closest similarity to near-natural counterparts with respect to biodiversity, ecosystem functioning and land cover characteristics indeed occurred for rewetted sites with annual median water tables close to the surface, we also observed sites with water tables close to the surface being strongly dissimilar to their near-natural counterparts. The general patterns indicate that restoration success depends on more than just a water table close to the surface (Supplementary Fig. 2).

**Locally novel ecosystems require functional understanding.**

With the existing dataset we cannot answer why some rewetted sites resemble near-natural counterparts right away, while others follow diverging trajectories for decades, indicating that the natural fens can become novel ecosystems sensu Hobbs et al.\(^1\), i.e., non-restorable at least for several decades, due to prolonged drainage (Fig. 4). Duration and intensity of drainage might likely matter here\(^3\) as these determine nutrient availability and irreversible changes in peat hydraulic variables (Fig. 4b). Existing knowledge from natural systems about ecological functions, external regulation, self-regulation, and ecosystem service provision appears hardly transferable for understanding and managing these locally novel ecosystems. Therefore, we call for a concerted action to assemble data about ecological functions and relevant meta information of rewetted peatlands as laid out by Bonnett et al.\(^4\). Understanding causes and consequences of restoration success will likely require the assessment of peat microbiomes and

**Fig. 2 Rewetted temperate fen peatland sites (open red) are more variable than near-natural sites (filled blue) and many rewetted sites are outside the near-natural range of variation.** Ordination (Non-metrical multidimensional scaling; NMDS) and analysis of similarity (ANOSIM) with its respective R value and p value provided in the figures. Dissimilarity is measured as Bray–Curtis distance for (a) vegetation based on 539 plant species (n\(_{\text{rewetted}}\) = 320, n\(_{\text{near-natural}}\) = 243), and Euclidean distance for (b) hydrology (based on annual median water table, annual minimum water table, annual maximum water table, annual amplitude of water table; n\(_{\text{rewetted}}\) = 320, n\(_{\text{near-natural}}\) = 243), (c) geochemistry (based on pH of pore water, electrical conductivity of the pore water in the top soil layer, bulk density, organic matter content of the top layer; n\(_{\text{rewetted}}\) = 16, n\(_{\text{near-natural}}\) = 47) and (d) land cover characteristics measured as 208 spectral-temporal metrics of high resolution optical Earth Observation data (n\(_{\text{rewetted}}\) = 258, n\(_{\text{near-natural}}\) = 114). The ellipses display 95%CI for each group. Sample size differs for the four response clusters according to data availability. The vegetation data are set into perspective against 10k vegetation plots randomly selected from > 90k vegetation plots from Europe of the displayed EUNIS-classes\(^2\) covering fen vegetation indicating the broad coverage of our near-natural sites. Note that no information about rewetting status is available for these background data. For vegetation, 63% of the rewetted sites are located outside the 95% confidence ellipse of the near-natural sites, for geochemistry 44%, for hydrology 20% and for spectral-temporal metrics 21%, respectively. Final stress of the ordinations are 0.16 for vegetation, 0.02 for hydrology, 0.04 for geochemistry, and 0.12 for land cover characteristics. The inserted boxplots display the median, quartiles (box), 1.5-fold quartile distance (whiskers) and extremes beyond the whiskers of all pairwise distances for rewetted sites in red and near-natural sites in blue (a and b, n\(_{\text{rewetted}}\) = 51040, n\(_{\text{near-natural}}\) = 29403; c, n\(_{\text{rewetted}}\) = 120, n\(_{\text{near-natural}}\) = 1081; d, n\(_{\text{rewetted}}\) = 33,135, n\(_{\text{near-natural}}\) = 6441). Significance of difference in mean values between rewetted and near-natural was evaluated by a permutation procedure as described in the methods section (**p < 0.001 for all four panels).
key microbial players in organic matter mineralization and GHG production and consumption and the GHG fluxes alongside vegetation, hydrology, pore water chemistry as well as chemical and physical peat properties.

Recently published studies showing that rewetted peatlands can act as carbon sinks even during extended dry and hot periods are encouraging, although the stop of continued carbon losses from peat degradation is the most pressing issue today. However, historical drainage of peatlands followed by rewetting creates locally novel ecosystems in many cases. Understanding and management schemes cannot be transferred from natural systems. Instead, an interdisciplinary, process-based understanding of the rewetted systems is urgently needed to prioritize, plan and implement restoration measures and to design their sustainable management. The UN Decade of Ecosystem Restoration starting just now is the critical period for achieving our commitments of the Paris Agreement on peat soils and for peatland science.

Methods

Site selection. A total of 320 rewetted sites was compared to 243 undrained, natural sites (Supplementary Data 1), covering the major distribution range of temperate fens across Europe (Fig. 1). We included all sites where local experts confirmed a dateable rewetting action, usually the blocking of drainage systems, which led to a mean annual water table of 25 cm below surface or higher. Rewetting occurred on average 9 years (min. 1 and max. 54 years) before data sampling. Land use before rewetting was agriculture (80% of all rewetted sites), forestry (10%) or peat extraction (10%). We acknowledge that few, if any, untouched and completely natural temperate fens exist nowadays in Europe. For our comparison, we used sites without direct drainage history, again relying on local expert knowledge. We confirmed this by carefully checking field conditions and aerial/satellite imagery for structures which would imply substantial former drainage and disregarded those with such features. Our study contains some of the least disturbed fens in temperate Europe (Eastern Poland and Belarus).

Response parameters. We included sites that provided data for at least two of the following four response clusters:

1. Vegetation: complete lists of vascular plants and bryophytes (539 species in total) based on 16 m² (median, ranging between 12 and 25 m²) with estimates of individual plant species cover. All vegetation data collections that we are confident in their power to estimate the temporal dynamics at the respective sites representatively. On average, 2.3 years were measured.

Fig. 3 Rewetted fen peatlands differ in relevant ecological parameters from near-natural fen peatlands. Comparison between rewetted (red) and near-natural (blue) sites for all parameters used in the analyses for hydrology (a–d) and for geochemistry (e–h) alongside selected other relevant parameters (i–m). ‘Tall helophyte cover sum’ (i) is the sum over all tall helophyte species cover values according to Chytry et al. Asterisks indicate significance of differences according to two-sided Wilcoxon-rank test for numerical and Chi² test for factorial response parameters (*p < 0.05; **p < 0.01; ***p < 0.001). n per group is provided as numbers next to the boxplots, n differs between parameters as data was not available for all sites and all parameters, respectively. The boxplots display the median, quartiles (box), 1.5-fold quartile distance (whiskers) and extremes beyond the whiskers.
and we included only data if available for at least one full year. All water table measurements have been made in direct vicinity to the vegetation relevé. We considered the annual median water table, annual minimum water table, annual amplitude of water table (max minus min) as relevant response parameters. Data for water table during the growing season were highly correlated to the annual water table data (annual median versus median throughout spring and autumn: \(r^2 = 0.89\); annual median versus median throughout autumn: \(r^2 = 0.94\); annual maximum versus median throughout spring and autumn: \(r^2 = 0.83\); annual maximum versus median throughout autumn: \(r^2 = 0.86\); annual minimum versus median throughout spring and autumn: \(r^2 = 0.91\); annual minimum versus median throughout winter and spring: \(r^2 = 0.73\).

(3) Geochemistry: pH and electrical conductivity of the pore water (0–60 cm), bulk density and organic matter content of the top soil layer (0–30 cm) sampled in summer for all sites included here alongside the vegetation data sampling. The parameters considered for geochemistry and also for hydrology had to be chosen in order to maximize data availability in parallel with meaningfulness. Note that, still, geochemical data are only available for a subset of the full data set (57 sites; see Fig. 3e–h for sample size per group). Despite the low sample size, it is important to note that the remaining sites cover almost the whole geographical range of the other response parameters (Eastern Poland, Eastern and Northern Germany, the Netherlands, Belgium, Wales) and no geographical or ecological bias was found. In addition, the remaining sites contain well-comparable pairs of rewetted and near-natural sites throughout this geographical space. While the selected parameters allow for relevant insights, more specific response parameters are required for process-based understanding of the biogeochemistry of rewetted peatlands.

(4) Land cover characteristics: spectral-temporal metrics for a full annual time series of Copernicus Sentinel-2 A/B data for 2018. The Sentinel-2 A/B constellation provides optical imagery of the Earth’s surface between 0.49–2.2 μm in ten spectral bands and at 10–20 m ground sampling distance at a theoretical acquisition frequency of 2.5–5 days. We here acquired all available Sentinel-2 A/B imagery for 2018 with cloud cover <70% from the ESA API Hub. We used all valid observations to derive spectral-temporal metrics from the time series. Spectral temporal metrics are statistical measures (e.g., average, minimum, maximum, quartiles, ...) per spectral band or index (e.g., MNDWI = (green – short wave infrared)/(green + short wave infrared)) using all available cloud- and shadow-free observations over time. The median count of clear-sky-observations per pixel across the sites is 45, while 90% of all sites featured 27 clear-sky observations or more. Both data processing to Analysis Ready Data as well as calculating spectral-temporal metrics was performed through the Framework for Operational Radiometric Correction for Environmental monitoring. Our analysis included data averaged over 3×3 pixels around the center plot location of the site. Different spatial aggregations (e.g., single pixels, 5×5 pixels around the center plot) led to highly similar results, implying that the intra-site variability was robust around locations of the vegetation survey. The inclusion of an annual series of Sentinel-1 synthetic aperture radar data (temporal metrics for VV and VH polarization, IW swath at 10 m resolution) for the same year did not affect the results.

The representativeness of our database is confirmed by the comparison to 10k vegetation plots from all over Europe. These have been randomly chosen from >90k vegetation plots of the European Vegetation Archive, classified as belonging to peatland types (Q2: “Poor fens”, Q4: “Calcareaous rich fens”, Q5: “Tall helophyte beds”, Qc: “Mires”) of the EUNIS habitat types and the checklist of bryophytes of Europe and harmonized to exclude nested taxa (option maxtaxtable = ‘AG1’ and ag = ‘conflict’ in function taxaval in R package vegdata).

Statistical analyses. An ordination (non-metrical multidimensional scaling, Bray-Curtis distance and step-across dissimilarities for those plot distances without common species with function ‘metaMDS’ from R-package ‘vegan’) was performed through the metaMDS of the vegan package. The ordination results imply that the response variables are suitable for the ordination analysis, however, a further discussion of the ordination results will be published in a separate study.
applied in parallel to an analysis of similarity (function ‘anosim’ from package ‘vegan’ 2.5-6.) to test for significance of differences among rewetted and near-natural peatlands. An Indicator Species Analysis function ‘indval’ from package ‘labds’ version 2.0-1) was applied to find plant species with significant affinity to rewetted or near-natural peatlands. NMDS ordinations and analyses of similarity based on Euclidean distance were calculated for the three other response clusters, i.e., hydrology, geochemistry, and land cover characteristics. Variance, quantified as pairwise distance in the response clusters, was compared between rewetted and near-natural sites by testing for significance of differences in mean values per group. As data points in pairwise dissimilarity matrices are not strictly independent and homoscedasticity and normal distribution of residuals was questionable, we relied on permutation based procedures for estimating the significance of differences in mean values of pairwise dissimilarities between groups (function ‘diffmean’ from package ‘simba’ version 0.3-5 with 1000 permutations per test).

In order to account for large variance in the predictors due to the broad spatial coverage of our database, additional analyses relied on pairwise comparisons between rewetted sites and near-natural counterparts. All sites were classified according to their biogeographic peatland region and hydrogenetic and ecological fen type according to Joosten et al. Then, the spatially closest near-natural sites by testing for significance of differences in mean values per group. As data points in pairwise dissimilarity matrices are not strictly independent and homoscedasticity and normal distribution of residuals was questionable, we relied on permutation based procedures for estimating the significance of differences in mean values of pairwise dissimilarities between groups (function ‘diffmean’ from package ‘simba’ version 0.3-5 with 1000 permutations per test). When assessing the significance of differences, we applied a Bonferroni correction for multiple comparisons.

To compare the extent of recovery in terms of microbial community composition, we used a non-metric multidimensional scaling (NMDS) on the Bray-Curtis dissimilarity of species occurrence patterns. The NMDS ordination was calculated and visualized using the package ‘vegan’ for R (Oksanen et al. 2018). To test for differences among sampling sites (rewetted, near-natural, and control) with respect to their microbial community composition, we performed an analysis of similarity (Anderson et al. 2008). The significance of the differences in microbial community composition among the sites was calculated using 999 permutations and the ‘adonis’ function from the ‘vegan’ package (Oksanen et al. 2018). For all statistical tests, a significance level of 0.05 was used. To assess the effect size of the differences, we calculated the pseudo-F statistic and the corresponding p-value and the Cohen’s d statistic (Hedges and Olkin 1981). The significance of differences in mean values per group. As data points in pairwise dissimilarity matrices are not strictly independent and homoscedasticity and normal distribution of residuals was questionable, we relied on permutation based procedures for estimating the significance of differences in mean values of pairwise dissimilarities between groups (function ‘diffmean’ from package ‘simba’ version 0.3-5 with 1000 permutations per test).

To assess the extent of recovery of the microbial community composition, we used a non-metric multidimensional scaling (NMDS) on the Bray-Curtis dissimilarity of species occurrence patterns. The NMDS ordination was calculated and visualized using the package ‘vegan’ for R (Oksanen et al. 2018). To test for differences among sampling sites (rewetted, near-natural, and control) with respect to their microbial community composition, we performed an analysis of similarity (Anderson et al. 2008). The significance of the differences in microbial community composition among the sites was calculated using 999 permutations and the ‘adonis’ function from the ‘vegan’ package (Oksanen et al. 2018). For all statistical tests, a significance level of 0.05 was used. To assess the effect size of the differences, we calculated the pseudo-F statistic and the corresponding p-value and the Cohen’s d statistic (Hedges and Olkin 1981).

Reporting summary

Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability

The data generated and analyzed in this study have been deposited in the Dryad database under https://doi.org/10.5061/dryad.08kprr532. The plot-based vegetation data used for comparison to this dataset is available at the European Vegetation Archive.

Code availability

All analyses were carried out with the functions and additional packages specified in the methods section in the free and open-source environment R.

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**Author contributions**

F.Ta., G.L. J.K. designed this study. A.K., B.L., C.A., E.S., E.V., G.L. F.Ta., H.J., L.C., J.K., L.K., M.W., N.W.M., R.v.D., T.U., W.-J.E., W.K. conceived the research question. A.K., A.S., A.P., B.T., C.A., C.O., C.S., D.Z., E.P., E.S., F.J., F.Ta., H.L., J.S., K.P., K.R., K.V., L.K., L.W., R.v.D., S.R., S.T., T.T., V.B., W.-J.E., W.K., Y.L. contributed data. F.Ta. assembled and harmonized the data. F.Ta. and D.M. checked all data and assigned the pairwise counterparts. F.J. harmonized the vegetation data with the additional vegetation plot data, added the EUNIS classification and ran the ordination for the vegetation data. S.v.R., F.S., F.Th. contributed the land cover characteristics data. J.K. analyzed the data and led the writing with all co-authors contributing.

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