



# Article (refereed) - postprint

van den Berg, Leon J.L.; Jones, Laurence; Sheppard, Lucy J.; Smart, Simon M.; Bobbink, Roland; Dise, Nancy B.; Ashmore, Mike R. 2016. **Evidence for differential effects of reduced and oxidised nitrogen deposition on vegetation independent of nitrogen load**.

#### © 2015 Elsevier Ltd.

This manuscript version is made available under the CC-BY-NC-ND 4.0 license <a href="http://creativecommons.org/licenses/by-nc-nd/4.0/">http://creativecommons.org/licenses/by-nc-nd/4.0/</a>

(CC) BY-NC-ND

This version available http://nora.nerc.ac.uk/512547/

NERC has developed NORA to enable users to access research outputs wholly or partially funded by NERC. Copyright and other rights for material on this site are retained by the rights owners. Users should read the terms and conditions of use of this material at <a href="http://nora.nerc.ac.uk/policies.html#access">http://nora.nerc.ac.uk/policies.html#access</a>

NOTICE: this is the author's version of a work that was accepted for publication in *Environmental Pollution*. Changes resulting from the publishing process, such as peer review, editing, corrections, structural formatting, and other quality control mechanisms may not be reflected in this document. Changes may have been made to this work since it was submitted for publication. A definitive version was subsequently published in *Environmental Pollution* (2016), 208 (B). 890-897. 10.1016/j.envpol.2015.09.017

www.elsevier.com/

Contact CEH NORA team at noraceh@ceh.ac.uk

The NERC and CEH trademarks and logos ('the Trademarks') are registered trademarks of NERC in the UK and other countries, and may not be used without the prior written consent of the Trademark owner.

- 1 This is an author-created version. The full article can be found at: Environmental Pollution
- 2 October 2015:DOI: 10.1016/j.envpol.2015.09.017

3

- 4 Evidence for differential effects of reduced and oxidised nitrogen deposition on
- 5 vegetation independent of nitrogen load
- 6 Leon JL van den Berg<sup>a,b,\*</sup>, Laurence Jones<sup>c</sup>, Lucy J Sheppard<sup>d</sup>, Simon M Smart<sup>e</sup>, Roland
- 7 Bobbink<sup>f</sup>, Nancy B Dise<sup>d</sup>, Mike R Ashmore<sup>g</sup>

8

9

- <sup>a</sup> Unie van Bosgroepen, Ede 6710AD, The Netherlands, l.vandenberg@bosgroepen.nl
- 11 b Radboud University Nijmegen, Nijmegen 6525AJ, The Netherlands
- 12 <sup>c</sup> Centre for Ecology & Hydrology, Environment Centre Wales, Bangor LL57 2UW, UK
- d Centre for Ecology & Hydrology, Bush Estate, Penicuik EH26 OQB, UK
- 14 <sup>e</sup> Centre for Ecology & Hydrology, Lancaster Environment Centre, Bailrigg, Lancaster LA1
- 15 4AP, UK
- <sup>f</sup> B-Ware research centre, Nijmegen 6503GB, The Netherlands
- 17 g Stockholm Environment Institute, University of York, Heslington, York YO10 5DD, UK

18

- 19 \*Corresponding author: Leon van den Berg
- 20 Unie van Bosgroepen, Ede 6710AD, The Netherlands, l.vandenberg@bosgroepen.nl

- 22 Capsule:
- 23 Effects of total N deposition and reduced and oxidised N deposition were studied across
- eight habitat types in the UK using data from the British Countryside Survey.
- 25 Highlights:
- 26 -N deposition was significantly related to species richness in all habitats except base-rich
- 27 mires.
- 28 -Form of N in deposition was related to biodiversity in grasslands and woodlands.
- 29 -Reduced N deposition was related to higher Ellenberg N values in all but one habitat type.

30 -Reduced N was negatively related to species richness in acid and mesotrophic grasslands.

#### Abstract

32

33 Nitrogen (N) deposition impacts natural and semi-natural ecosystems globally. The 34 responses of vegetation to N deposition may, however, differ strongly between habitats and 35 may be mediated by the form of N. Although much attention has been focused on the 36 impact of total N deposition, the effects of reduced and oxidised N, independent of the total 37 N deposition, have received less attention. In this paper, we present new analyses of national monitoring data in the UK to provide an extensive evaluation of whether there are 38 differences in the effects of reduced and oxidised N deposition across eight habitat types 39 (acid, calcareous and mesotrophic grasslands, upland and lowland heaths, bogs and mires, 40 base-rich mires, woodlands). We analysed data from 6860 plots in the British Countryside 41 Survey 2007 for effects of total N deposition and N form on species richness, Ellenberg N 42 values and grass:forb ratio. Our results provide clear evidence that that N deposition affects 43 species richness in all habitats except base-rich mires, after factoring out correlated 44 45 explanatory variables (climate and sulphur deposition). In addition, the form of N in deposition appears important for the biodiversity of grasslands and woodlands but not 46 47 mires and heaths. Ellenberg N increased more in relation to NH<sub>x</sub> deposition than NO<sub>y</sub> deposition in all but one habitat type. Relationships between species richness and N form 48 49 were habitat-specific: acid and mesotrophic grasslands appear more sensitive to NH<sub>x</sub> deposition while calcareous grasslands and woodlands appeared more responsive to NO<sub>v</sub> 50 deposition. These relationships are likely driven by the preferences of the component plant 51 species for oxidised or reduced forms of N, rather than by soil acidification. 52 53 54 55 56 57

#### Keywords

58

59

60

61

NH<sub>x</sub>:NO<sub>y</sub> ratio, N deposition, countryside survey, acidification, grassland, heathland, bogs

#### Introduction

There is widespread evidence across the globe, from both experiments and field surveys, of the significant ecological impacts of nitrogen (N) deposition on semi-natural ecosystems of low nutrient status (e.g. Bobbink et al. 2010), which also carries economic costs (Jones et al. 2014). However, interpretation and quantification of these effects, and predictions of the benefits of emission control policies, need to consider the different components of N deposition (Brink et al. 2011). There are two main chemical forms - reduced N (ammonia, NH<sub>3</sub> and ammonium, NH<sub>4</sub><sup>+</sup>) emitted primarily from agricultural sources, and oxidised N (nitrogen oxides, NO<sub>v</sub>, nitric acid, HNO<sub>3</sub><sup>-</sup> and nitrate NO<sub>3</sub><sup>-</sup>) emitted primarily from fossil fuel combustion. In addition, N deposition may be in the form of dry deposition of gases and aerosols, which is most important close to sources, and in regions of the world with low rainfall, and as wet deposition as snow, dew, cloud or rainwater, which are important in more remote regions and in areas with high rainfall.

The mechanisms underlying the ecological effects of N deposition include direct toxicity, growth stimulation and competitive exclusion, soil acidification and increased susceptibility to other abiotic and biotic stresses (e.g. Bobbink et al. 1998, Roem and Berendse 2000). There are strong reasons, which have been recently reviewed by (Stevens et al. 2011), for expecting that there may be different effects of reduced and oxidised N deposition for each of these mechanisms. For example, foliar uptake of gaseous NH<sub>3</sub> is more likely to be directly toxic than uptake of gaseous nitrogen oxides, while soil NH<sub>4</sub><sup>+</sup> is more likely to be toxic to plant roots than soil NO<sub>3</sub><sup>-</sup> (Sheppard et al. 2011, Sheppard et al. 2014). Plant species also differ strongly in their preference and tolerance for NH<sub>4</sub><sup>+</sup> or NO<sub>3</sub><sup>-</sup> uptake from soil solution with species of acidic habitats generally more tolerant of higher soil ammonium (Falkengrengrerup and Lakkenborgkristensen 1994). The soil NH<sub>4</sub><sup>+</sup>/NO<sub>3</sub><sup>-</sup> ratio is partly a function of the ratio in atmospheric deposition, but also of the degree of nitrification in soils; high rates of nitrification result in a lower soil solution NH<sub>4</sub><sup>+</sup>/NO<sub>3</sub><sup>-</sup> ratio, which may reduce the risk of direct NH<sub>4</sub><sup>+</sup> toxicity but may increase acidification because of the greater oxidation to NO<sub>3</sub><sup>-</sup>.

Experimental studies provide some evidence of the differential effects of reduced and oxidised N deposition. For example, van den Berg et al. (2008) showed that higher NH<sub>4</sub><sup>+</sup>/NO<sub>3</sub><sup>-</sup> ratios in deposition to heathland mesocosms had significant adverse effects on acid-sensitive species but not on acid-tolerant species that were also tolerant of high soil NH<sub>4</sub><sup>+</sup>/NO<sub>3</sub><sup>-</sup> ratios. This effect was lost in limed mesocosms, suggesting that acidification at higher NH<sub>4</sub><sup>+</sup>/NO<sub>3</sub><sup>-</sup> ratios was the key driving mechanism. By contrast, in Mediterranean maquis vegetation, the application of both NH<sub>4</sub><sup>+</sup> and NO<sub>3</sub><sup>-</sup> increased biomass but not plant diversity, while NH<sub>4</sub><sup>+</sup> alone increased plant diversity but not biomass (Dias et al. 2014); these effects can at least partly be explained by the different responses of individual species to total N inputs or to reduced N deposition specifically.

A combination of targeted field surveys and analysis of nationwide surveillance data over 40 the last decade have provided a strong body of evidence of the impacts of N deposition. 41 Strong negative associations between N deposition and species richness have been reported 42 in acid grasslands (Stevens et al. 2004, Duprè et al. 2010, Stevens et al. 2010a), grasslands, 43 44 heathlands and bogs (Maskell et al. 2010, Caporn et al. 2014, Field et al. 2014)and sand 45 dunes (Jones et al. 2004). In acid grasslands, this negative association is linked to declines in forb species richness and a corresponding increase in graminoids (Maskell et al. 2010) with 46 47 differential responses of individual forb species to N deposition (Payne et al. 2013). In acid grasslands acidification rather than eutrophication may be the main driver of change 48 49 (Stevens et al. 2010b), but the relative influence of sulphur versus nitrogen as the driver of 50 acidification has not been separated.

However, in some other habitats; for example in calcareous grasslands, gradient surveys have shown no significant association between N deposition and species richness (Maskell et al. 2010). However, high rates of N deposition have been associated in calcareous grassland plots with an increase in grass:forb ratio (Maskell et al. 2010) and a decline in species diversity and in the frequency of characteristic species (van den Berg et al. 2011). This latter study suggests that, while direct effects of N deposition were responsible for shifts in diversity, effects on herb species number reflect indirect effects of both N and S deposition on soil acidity.

51

52

53

54

55

56 57

58

59

60

61

62

63

64

65

66

67

68

69 70

71

72

73

74

75 76

77 78 These and other findings from field surveys suggest that the responses to N deposition of vegetation characteristics in different habitats may be at least partly explained by differences in the underlying mechanisms of impact of reduced and oxidised N, mediated by soil pH, with acidification effects prevailing in poorly-buffered habitats and eutrophication effects in well-buffered habitats. Few field surveys have tried to separately evaluate the strength of associations with reduced and oxidised nitrogen but were only able to do so with relatively low number of samples/sites (Caporn et al. 2014, Field et al. 2014). Three studies have showed adverse changes in vegetation composition that were significantly correlated with reduced N deposition but not with oxidised N deposition: an increase in mean Ellenberg fertility index in semi-natural grassland and heaths/bogs between 1990 and 1998 in UK Countryside Survey data (Smart et al. 2004); a loss of species with a low Ellenberg fertility index in UK national recording data between 1987 and 1999 (McClean et al. 2011); and increases in graminoid cover and decreases in lichen cover in heathlands (Southon et al. 2013). A further study showed effects only of dry deposition of NH<sub>x</sub> and no effect of wet reduced or oxidised N on abundance of N sensitive epiphytic lichens (Seed et al. 2013).

However, interpretation of such field surveys is difficult due to problems of spatial autocorrelation, and the confounding effects of other environmental and land use changes. The levels of reduced and oxidised N deposition are often highly correlated (areas of low reduced N usually have low oxidised N, etc); in addition, the range and spatial variability of

reduced N deposition is often greater than that of oxidised N deposition, thereby increasing the probability of detecting a statistically significant association with vegetation characteristics (e.g. Smart et al. 2012). In this paper, we present new analyses of national surveillance data in the UK to provide a more rigorous evaluation of whether there are differences in the effects of reduced and oxidised N deposition that are more robust to statistical limitations. The data that are used here provide a greater sample size and spatial scope that includes almost the complete N deposition range in the UK and allows us to evaluate our mechanistic understanding of the differential effects of the two forms of N deposition in different habitats and on different groups of species. In our analysis we focus on species richness of vascular plants as a measure of biodiversity, Ellenberg N as a measure of nutrient status (Diekmann and Falkengren-Grerup 2002) and grass:forb ratio as a measure of competitive dominance effects. We hypothesise that:

- -The form of N (oxidised NO<sub>y</sub>, or reduced NH<sub>x</sub>) in deposition has an effect on vegetation composition that is independent of, and additional to, that of total N deposition.
- -Reduced N deposition has a greater impact on vegetation composition than oxidised Ndeposition.

### 1 Methods

### 2 Vegetation data

- 3 The effect of N deposition on vegetation was assessed using vegetation data obtained from
- 4 6860 plots (2x2m) from the UK Countryside Survey 2007 (Carey et al. 2008). For each plot,
- 5 total species richness, grass:forb ratio and mean Ellenberg N values were calculated. Species
- 6 richness was defined as the sum of all vascular plants in each 2x2 m plot. Grass to forb ratio
- 7 was based on the cover of grass species (*Poaceae*) divided by the cover of forb species.
- 8 Cover-weighted average Ellenberg N numbers (Ellenberg et al. 1991) that were modified for
- 9 the UK (Hill et al. 2004) were calculated based on the cover per 2x2m plot to obtain strong
- correlates of species responses to nutrient availability and succession (Vile et al. 2006).
- 11 The vegetation in each plot was classified according to the UK National Vegetation
- 12 Classification (NVC) and plots were pooled into broad groups of similar habitat (see table in
- supplementary material). Earlier studies have shown that soil pH or base saturation can
- explain species richness and can affect the responses of the vegetation and the ecosystem
- to N deposition (van den Berg et al. 2005, Stevens et al. 2006). Therefore, habitats that
- belong to a similar broad NVC classification but differ strongly in average soil pH and/or
- base cation content (heaths, mires and grasslands) were subdivided according to pH for
- analysis. The resulting broad habitat types in this study were: bogs and mires (acidic), base
- 19 rich mires and fens, upland dry heaths, lowland dry heaths, calcicolous grasslands,
- 20 mesotrophic grasslands, calcifugous grasslands and woodlands. Bogs and mires comprise
- 21 the NVC classes: M1 to M21 (mires) and H3, H4 and H5 (wet heaths). Base rich mires and
- fen habitat consist of the NVC classes M22 to M38. Upland dry heaths are NVC classes H10
- to H22, lowland dry heaths are H1, H2, H6 to H9, calcicolous grasslands are CG1 to CG14,
- 24 mesotrophic grasslands are MG1 to MG13, calcifugous grasslands are U1 to U21 and
- woodlands are W1 to W25. All sub-communities were included.

### 26 Atmospheric deposition and climatic data

- 27 Climatic factors such as precipitation and temperature were included in our models as these
- are known to affect species richness (Cleland et al. 2013). Average annual temperature (°C)
- and average annual precipitation (mm), calculated over a 5 year period 2000-2005 were
- 30 obtained from UK Meteorological Office (www.metoffice.co.uk). Sulphur (S) deposition, that
- 31 peaked in the UK in the 1970s can have an acidifying effect on the soil (Kirk et al. 2010) and
- may thereby affect species richness (McGovern et al. 2011). Historical data on S deposition
- was therefore included in our models to account for potential legacy of soil acidification
- effects due to sulphur. Modelled N deposition data for each plot were obtained from the
- 35 Centre of Ecology and Hydrology (CEH) for the year 2007; data from 1987 were used for
- 36 historical S deposition. Climate and pollution data were all at 5x5km resolution. Total N
- deposition ranged from 5.1 to 54.2 kg N ha<sup>-1</sup>yr<sup>-1</sup> while S deposition ranged from 5.0 to 43.5
- 38 kg S ha<sup>-1</sup>yr<sup>-1</sup>. Oxidised and reduced N were included in our models as the sum of wet and dry

- NO<sub>V</sub> or NH<sub>X</sub> deposition and expressed in kg N ha<sup>-1</sup>yr<sup>-1</sup>. NO<sub>V</sub> deposition ranged from 2.5 to
- 40 25.6 kg N ha<sup>-1</sup>yr<sup>-1</sup>, NH<sub>x</sub> deposition ranged from 2.3 to 36.1 kg N ha<sup>-1</sup>yr<sup>-1</sup>. The ranges of N and
- S deposition for each habitat are different and depend on their geographical distribution.
- 42 Linear models
- 43 Multicolinearity is common between variables such as N deposition and the different forms
- of N in deposition, S deposition and climatic variables. In our dataset, NH<sub>x</sub> and NO<sub>y</sub>
- deposition were highly correlated (r=0.69, p<0.001) and could therefore not be analysed
- simultaneously. In addition, total N deposition and either NH<sub>x</sub> or NO<sub>y</sub> deposition were highly
- 47 correlated (r=0.95, p<0.001 and r=0.89, p<0.001 respectively). Analysis to determine the
- 48 effect of N form was therefore performed using linear models taking two different
- 49 approaches that each overcome problems typically associated with multicolinearity and that
- each test specific hypotheses. Linearity of the relationship between the predictor factors
- and the dependent variables were tested in single linear regressions. If needed, data were
- transformed to meet assumptions of linearity.
- In the first method (models coded with A), the effect of NH<sub>x</sub>/NO<sub>y</sub> ratio in deposition was
- tested against the effects of total N deposition. For this analysis, multiple regressions were
- 55 performed with dependent variables: species richness, cover weighted Ellenberg N
- 56 (hereafter Ellenberg N) and grass:forb ratio that were regressed on the explanatory
- variables: total N deposition, NH<sub>x</sub>/NO<sub>y</sub> ratio, S deposition, precipitation and temperature. All
- 58 models were at first explicitly tested for spatial autocorrelation in the response variable and
- residuals by inspection of semi-variograms using generalized linear mixed-effect models
- 60 (GLMM). In these models, a correlation structure was added to correct for spatial
- autocorrelation. Correlation structures such as corExp, and corSpher were used with the
- 62 "form=~Easting+Northing" argument in the correlation option to calculate the Euclidean
- distances (using Pythagoras theorem) between sites with coordinates given by Easting and
- Northing. When spatial autocorrelation was not present or not severe, multiple linear
- regression models were used.
- In the second method (models coded with B), the additional effects of either NH<sub>x</sub> or of NO<sub>y</sub>
- on species richness, Ellenberg N and grass:forb ratio were tested after taking into account
- the variation explained by S deposition, precipitation, temperature and the other form of N.
- 69 In this analysis, the residuals of a model that regresses a predictor against NH<sub>x</sub>, S deposition,
- 70 precipitation and temperature were regressed in a second model against NO<sub>v</sub>. i.e. the
- 71 relationship between NO<sub>y</sub> deposition and the unexplained variance of the model was tested.
- 72 The calculation was then repeated for an analysis of NH<sub>x</sub> on the residuals of a model that
- included NO<sub>y</sub>. Given that the data cover a substantial range of NO<sub>y</sub> and NH<sub>x</sub> deposition, and
- making the assumption that the observed responses with N deposition are linear, any
- differential effect of reduced and oxidised N deposition is independent of the range of the
- length of the deposition gradient. In this way, the slope coefficients (effect sizes) that are

78 accounting for other sources of (co-correlated) variation. 79 80 Multicolinearity in the models was detected by calculating the Pearson correlation coefficient among pairs of the predictors and by calculating the variance inflation factors 81 82 (VIF) for each predictor in the model. Predictors with VIF of less than 4 were maintained in the models since these indicate that problems with multicolinearity are not severe (Gujarati 83 84 1995). Predictor variables that were highly correlated (VIF>4) were not analysed in the same 85 model. In an additional step, multicolinearity was explored by comparing the beta-86 coefficients of the explanatory variables that were obtained in a multiple regression with the beta coefficients from single regressions of these explanatory variables. In this analysis, 87 88 major changes in beta coefficient or changes in sign indicate multicolinearity between explanatory variables that needs to be accounted for. 89 90 Statistical analysis were performed using the 'nlme' package in the 'R' (version 2.9.0) 91 statistical and programming environment (R Development Core Team 2008) and SPSS 92 version 21 (IBM statistics). 93

derived allow a comparison of the independent effects of either NH<sub>x</sub> and NO<sub>y</sub>, after

#### 1 Results

- 2 N deposition and  $NH_x/NO_y$  ratio
- 3 Our analysis shows that species richness was negatively affected by total N deposition for all
- 4 habitats apart from base rich mires (no significant effect) and calcareous grasslands (a
- 5 significant positive effect) (Table 1 and 2; Figure 1). Coefficients were comparable for the
- 6 habitats mesotrophic grasslands, bogs and mires, woodlands, acidic grasslands and dry
- 7 upland heaths. The strongest negative coefficient was found for dry lowland heath.
- 8 Species richness of all three grassland habitats was negatively related to NH<sub>x</sub>/NO<sub>y</sub> ratio in
- 9 deposition when effects of total N deposition were accounted for (Table 1 and 2; Figure 2).
- 10 For woodlands a positive relationship between species richness and NH<sub>x</sub>/NO<sub>y</sub> ratio was
- found, while there was no significant effect of N form on the upland and lowland heathlands
- and the base-rich mires, bogs and mires. Analyses of residuals against NH<sub>x</sub> and NO<sub>y</sub> (Models
- B) in all cases were consistent with responses shown by NH<sub>x</sub>/NO<sub>y</sub> ratio (Models A). These
- analyses showed that the negative effects on species richness in all three grassland habitats
- were driven by strong negative effects of NH<sub>x</sub>. Species richness in calcareous grasslands was
- also positively related to NO<sub>y</sub> (Table 2). In woodlands in contrast, the negative effects on
- species richness were strongly related to NO<sub>y</sub> deposition.
- 18 Total N deposition increased Ellenberg fertility index for bogs & mires, base rich mires,
- mesotrophic grasslands and calcareous grasslands, but decreased fertility index in dry
- 20 lowland heath and acidic grasslands (Table 1 and 2; Figure 3). There was no significant effect
- 21 on fertility index in upland heaths or woodlands. In all habitats apart from the base rich
- 22 mires, there was a significant positive relationship of comparable size between NH<sub>x</sub>/NO<sub>y</sub>
- ratio and the Ellenberg fertility index (Figure 4). However, the form of N responsible and the
- 24 nature of the relationship differed among the habitats. For dry lowland heath and acidic
- 25 grassland, this ratio effect was driven by a strong negative relationship with oxidised N, i.e.
- 26 NO<sub>y</sub> reduced fertility index. In the case of the bogs & mires, mesotrophic grasslands and
- 27 woodlands, the ratio effect was caused by a positive relationship with reduced N, i.e. NH<sub>x</sub>
- 28 increased fertility index. For calcareous grassland there was both a negative relationship for
- 29 oxidised N and a positive relationship for reduced N with Ellenberg fertility scores.
- 30 Grass: forb ratios increased with greater N deposition in upland and lowland heathland, bogs
- 8 mires and acidic grasslands (Figure 5). Only in calcareous grasslands, grass:forb ratio was
- found to be lower with increased N deposition. There was no effect on grass:forb ratio in
- base rich mires, mesotrophic grasslands or woodlands. In acidic and calcareous grasslands
- 34 the increased grass:forb ratio was associated with increased NH<sub>x</sub>/NO<sub>y</sub> ratio (Table 2, Figure
- 35 6), but separate relationships for either reduced N or oxidised N were not significant and
- could not be used to infer which form of N was more responsible.
- 37 Additional environmental factors

Climate variables were frequently a significant explanatory variable for total species richness and Ellenberg fertility score (Table 1 and 2). Precipitation was negatively associated with species richness in the acidic and calcareous grasslands but positively associated with species richness in mesotrophic grasslands, dry upland heath and bogs & mires. A higher precipitation was associated with a lower Ellenberg fertility score in most habitats.

Temperature was positively associated with Ellenberg fertility scores. Past sulphur deposition also showed some significant effects. Sulphur deposition showed a significant relationship with species richness in acidic grasslands (negative), Ellenberg fertility scores for mesotrophic and calcareous grasslands (positive) and grass:forb ratio for bogs & mires (negative). This highlights the importance of factoring out these co-correlated variables to genuinely extract any relationships due to N deposition or N form.

#### Discussion

1

3

4

5 6

7

8

9

10

11 12

13 14

15

16 17

18 19

20

21

22

23

2425

2627

28 29

## 2 Total N deposition effects on species richness

Almost all habitats showed a negative relationship between total N deposition and species richness, largely corroborating previous gradient studies for acid grasslands, heathlands and bogs (Stevens et al. 2004, Duprè et al. 2010, Henrys et al. 2011, Caporn et al. 2014). In a previous study using 1998 Countryside Survey data, as opposed to the 2007 data used in our analysis, Maskell et al. (2010) found significant negative relationships for acid grassland and heathlands, but not for calcareous or mesotrophic grassland, a pattern consistent with our findings. Although woodlands showed a negative relationship for species richness in this study, Verheyen et al. (2012) suggest that species richness changes in woodlands are more attributable to management than to N deposition. The positive relationship in calcareous grasslands runs contrary to findings in most other habitats and may reflect differences in the types of grassland included in this category that are due to a combination of glacial history, biogeographical regions, altitude and management. The calcareous grasslands include the species rich CG1 and CG2 and the relative species poor CG10 and CG11. These habitat types are different with respect to species numbers, management and altitude. Separate analysis of the most abundant communities within the dataset (95% of the calcareous grassland records), the relatively species rich communities (UK NVC classes CG1 and CG2) and relatively species-poor communities (CG10 and CG11) showed no significant relationship with N deposition in either case, which is in line with other surveys in calcareous grasslands showing no effect of N (Bennie et al. 2006, van den Berg et al. 2011) and similar to Maskell et al (2010) who used a subset of NVC classes (CG2,3,4,6,8,10,11). A lack of significant relationships with N deposition may be caused by differences in local

A lack of significant relationships with N deposition may be caused by differences in local management that is aimed specifically at the conservation of high species diversity and in which grazing regimes are implemented to prevent the dominance of eutrophic species. In this study, local management was not taken into account. Base rich mires showed no relationship, but there are no other studies in this habitat against which to compare a response. However, negative effects of N on species richness have been shown in calcareous dune grasslands (Jones et al. 2004, Field et al. *in press*), suggesting that base-rich habitats are not immune to N impacts.

303132

33 34

35 36

37

38

39

40

The decline in species richness was accompanied by an increased grass:forb ratio in the acidic, open habitats (acidic grasslands, dry upland and lowland heaths and bogs and mires). In contrast, the highly buffered alkaline habitats base rich mires and calcareous grasslands, showed no or even a negative relationship between grass:forb ratio and N deposition. These results are in agreement with previous studies on acidic habitats that showed an increased grass:forb ratio with increasing N deposition due to either loss of forb species richness (Maskell et al. 2010, Payne et al. 2013) or increased grass encroachment (Remke et al. 2009, Friedrich et al. 2011, Provoost et al. 2011). Grass encroachment and a decline in forb species in acidic ecosystems are often attributed to accelerated acidification of the soil leading to a

depletion of base cations and increased availability of potential toxic metals such as iron and aluminium (De Graaf et al. 1997, Horswill et al. 2008). Both (historical) deposition of sulphur and N deposition are known causes for acidification (RoTAP 2012) and N deposition also results in eutrophication. However, our data does not allow us to disentangle the effects of eutrophication and acidification due to N deposition. Note that grass:forb ratio only increased in 4 of the 8 habitats, and therefore is not a consistent indicator of N impact.

 et al. 2014).

- Base rich habitats and, to a lesser extent, bogs and mires increased in Ellenberg fertility index with increasing N deposition which was not necessarily accompanied with a loss in species richness suggesting that elevated N deposition in these habitats results in a shift in species composition favouring more nutrient-loving species. In contrast, acid grasslands and lowland heaths show a small decline in fertility index with increasing N deposition. Others have also reported lower Ellenberg N values with higher N deposition in acid grasslands (Maskell et al. 2010) and heathlands (Caporn et al. 2014) and these relationships may be linked to the high correlation between Ellenberg N and Ellenberg R, suggesting mechanisms such as acidification to operate in these systems. The exact mechanisms for these relationships are however not known and need further exploration at the species level of both vascular plants and bryophytes; studies have shown much greater effects of N deposition on bryophyte species richness than vascular plant species richness (e.g. Caporn
- 61 N form and the relative influence of reduced versus oxidised N
  - NH<sub>x</sub> deposition and NO<sub>y</sub> deposition are highly correlated. In addition, NH<sub>x</sub> was highly correlated to total N deposition. Separate analysis of NH<sub>x</sub> and NO<sub>y</sub> effects in models that allowed us to factor out the variance that was explained by either one of the N forms was therefore considered the best method to compare effects of these N forms, after taking account of other variables and the multicolinearity that existed in the datasets. Since both forms are correlated the variance that is explained by one N form, and which is factored out, is likely to contain some degree of variance that in fact should be attributed to the other form. The method that we employed here is therefore considered conservative in its estimation of effect sizes and significance levels.
  - The range of  $NH_x$  (2.3 36.1 kgNha<sup>-1</sup>y<sup>-1</sup>) exceeds that of  $NO_y$  (2.5 25.6 kgNha<sup>-1</sup>y<sup>-1</sup>) over all habitats together. Although the gradient length of explanatory variables may affect the outcome of the analysis in small data sets (Smart and Scott 2004), large datasets such as the CS data capture a good proportion of the relationship (i.e. not just a small segment), even with smaller ranges of the explanatory variable. Our analysis is based on the assumption that the relationships are linear (transformations were applied when necessary) between the response variable and either  $NH_x$  or  $NO_y$  and estimations of the effect sizes of the relationships are therefore considered relatively unaffected by the length of the gradients. Plots of beta coefficients of the regressions for  $NO_y$  and  $NH_x$  against N-gradient length

confirm that the beta coefficients were indeed not affected by gradient length in the N ranges that we tested (data not shown). In addition, the modelled  $NH_x$  data may be more prone to error in predicting the actual  $NH_x$  deposition at each site. Although this increased scatter reduces the likelihood of finding a significant relationship with  $NH_x$ , the longer gradient length partly offsets this problem. Therefore the effect sizes give a good indication of the relative influence of  $NH_x$  or  $NO_y$ .

The form of N in deposition independent of N load did affect species richness, but only in certain habitats. N form altered species richness in all grasslands and in woodlands but not in the mires and heaths. The lack of response of species richness and also grass:forb ratio, to N form in heaths and mires may be due to the prevailing acidic conditions, restraining nitrification rates with naturally high soil  $NH_4^+/NO_3^-$  ratios and low base cation concentrations (e.g. De Graaf et al. 2009). Many species of acidic habitats, such as ericoids, are generally adapted to elevated  $NH_4^+$  concentrations and tolerate high  $NH_4^+$  concentrations (De Graaf et al. 1998, Britto and Kronzucker 2002, Sheppard et al. 2014).

Where N form was important, in the acidic and mesotrophic grasslands NH<sub>x</sub> appeared to be more important than NO<sub>y</sub> as a driver of species richness decline, corroborating an experimental study on N form in acid grasslands (Dorland et al. 2013). However, in the woodlands and in calcareous grassland, NO<sub>x</sub> was more important than NH<sub>y</sub>, having a positive effect on species richness in the grassland but a negative effect in the woodland. Nitrification and mineralisation in woodlands can be very high (Falkengrengrerup and Lakkenborgkristensen 1994, Falkengren et al. 1998). Atmospheric deposition of oxidised N may therefore favour nitrophilous species such as bramble and nettle that outcompete slow-growing forb and shrub species that are more adapted to ammonium nutrition (such as *Vaccinium myrtillus*), corroborating a recent simulation study (Stevens et al. this volume). The positive impact of NO<sub>y</sub> on calcareous grassland species richness may relate to the preference of many calcareous species for available N in oxidised rather than in reduced form.

The question remains why the species richness and composition of only some habitats are sensitive to N form, even though the fertility index of almost all habitats responded to N form. N form may alter species composition through preferences of the component species for oxidised or reduced N, through direct toxicity of NH<sub>3</sub> and NH<sub>4</sub><sup>+</sup> (Britto and Kronzucker 2002, van den Berg et al. 2005, Sheppard et al. 2011), or through indirect effects mediated by N-induced acidification (e.g. Bobbink et al. 1998, Stevens et al. 2011), which would be more apparent in acidic habitats. The lack of significance of N form in the more acidic habitats suggests that acidification is not the main cause. However, experimental studies have shown that elevated NH<sub>4</sub><sup>+</sup>/NO<sub>3</sub><sup>-</sup> ratios in deposition result in a decline of acid-sensitive species but not of acid-loving species tolerant of high soil NH<sub>4</sub><sup>+</sup>/NO<sub>3</sub><sup>-</sup> ratios (Paulissen et al.

2004, van den Berg et al. 2008). The response in acidic and mesotrophic grasslands may therefore reflect the abundance of species that are sensitive to reduced N in these neutral to moderately acidic habitats compared with the more strongly acidophile vegetation in heaths, bogs and mires. Clearly responses to N form are habitat-specific, and may be driven by the preference or tolerance of the component species for N in oxidised or reduced forms.

In conclusion, this study has shown that N affects species richness in almost all habitats, after correlating factors such as temperature, rainfall and historical sulphur deposition have been factored out. The form of N is important, with fertility index increasing with NH<sub>x</sub>/NO<sub>y</sub> ratio in almost all habitats. However, the effects of the ratio on species richness were only found in certain habitats (grasslands and woodland), not in others (mires and heaths). In habitats where there were differential effects of one N form or the other, acidic and mesotrophic grassland were more sensitive to NH<sub>y</sub>, while calcareous grassland and woodland were more sensitive to NO<sub>x</sub>. This study suggests that, contrary to our original hypothesis, sensitivity to N form is more likely due to the inherent preferences of component species for oxidised or reduced N, rather than linked to soil acidification. However, those preferences are related to soil pH with NH<sub>4</sub>-loving species generally more prevalent on acidic soils.

### Acknowledgements

We thank Lindsay Maskell for her help with the CS-data and David Fowler for making the N deposition data available.

#### References

- Bennie, J., M. O. Hill, R. Baxter, and B. Huntley. 2006. Influence of slope and aspect on long-term vegetation change in British chalk grasslands. Journal of Ecology **94**:355-368.
- Bobbink, R., K. Hicks, J. Galloway, T. Spranger, R. Alkemade, M. Ashmore, M. Bustamante, S.
   Cinderby, E. Davidson, F. Dentener, B. Emmett, J. W. Erisman, M. Fenn, F. Gilliam, A. Nordin,
   L. Pardo, and W. De Vries. 2010. Global assessment of nitrogen deposition effects on
   terrestrial plant diversity: a synthesis. Ecological Applications 20:30-59.
  - Bobbink, R., M. Hornung, and J. G. M. Roelofs. 1998. The effects of air-borne nitrogen pollutants on species diversity in natural and semi-natural European vegetation. Journal of Ecology **86**:717-738.
- Brink, C., H. van Grinsven, B. H. Jacobsen, A. Rabl, I.-M. Gren, M. Holland, Z. Klimont, K. Hicks, R.
  Brouwer, R. Dickens, J. Willems, M. Termansen, G. Velthof, R. Alkemade, M. van Oorschot, and J. Webb. 2011. Costs and benefits of nitrogen in the environment. Pages 513-540 *in* M.
  A. Sutton, C. M. Howard, J. W. Erisman, G. Billen, A. Bleeker, P. Grennfelt, H. van Grinsven, and B. Grizzetti, editors. The European nitrogen assessment. Cambridge University Press, Cambridge.
  - Britto, D. T. and H. J. Kronzucker. 2002. NH4+ toxicity in higher plants: a critical review. Journal of Plant Physiology **159**:567-584.
    - Caporn, S. J. M., J. A. Carroll, N. B. Dise, and R. J. Payne. 2014. Impacts and indicators of nitrogen deposition in moorlands: Results from a national pollution gradient study. Ecological Indicators **45**:227-234.
    - Carey, P. D., S. Wallis, P. M. Chamberlain, A. Cooper, B. A. Emmett, L. C. Maskell, T. McCann, J. Murphy, L. R. Norton, B. Reynolds, W. A. Scott, I. C. Simpson, S. M. Smart, and J. M. Ullyett. 2008. Countryside Survey: UK Results from 2007. NERC/Centre for Ecology & Hydrology.
    - Cleland, E. E., S. L. Collins, T. L. Dickson, E. C. Farrer, K. L. Gross, L. A. Gherardi, L. M. Hallett, R. J. Hobbs, J. S. Hsu, L. Turnbull, and K. N. Suding. 2013. Sensitivity of grassland plant community composition to spatial vs. temporal variation in precipitation. Ecology **94**:1687-1696.
    - De Graaf, M. C. C., R. Bobbink, J. G. M. Roelofs, and P. J. M. Verbeek. 1998. Differential effects of ammonium and nitrate on three heathland species. Plant Ecology **135**:185-196.
    - De Graaf, M. C. C., R. Bobbink, N. A. C. Smits, R. Van Diggelen, and J. G. M. Roelofs. 2009. Biodiversity, vegetation gradients and key biogeochemical processes in the heathland landscape. Biological Conservation **142**:2191-2201.
    - De Graaf, M. C. C., R. Bobbink, P. J. M. Verbeek, and J. G. M. Roelofs. 1997. Aluminium toxicity and tolerance in three heathland species. Water Air and Soil Pollution **98**:229-239.
    - Dias, T., A. Clemente, M. A. Martins-Loucao, L. Sheppard, R. Bobbink, and C. Cruz. 2014. Ammonium as a Driving Force of Plant Diversity and Ecosystem Functioning: Observations Based on 5 Years' Manipulation of N Dose and Form in a Mediterranean Ecosystem. Plos One 9.
  - Diekmann, M. and U. Falkengren-Grerup. 2002. Prediction of species response to atmospheric nitrogen deposition by means of ecological measures and life history traits. Journal of Ecology **90**:108-120.
- Dorland, E., C. J. Stevens, C. Gaudnik, E. Corcket, S. Rotthier, K. Wotherspoon, M. Jokerud, V.
  Vandvik, M. B. Soons, M. M. Hefting, P. A. Aarrestad, D. Alard, M. Diekmann, C. Dupre, N. B.
  Dise, D. J. G. Gowing, and R. Bobbink. 2013. Differential Effects of Oxidised and Reduced
  Nitrogen on Vegetation and Soil Chemistry of Species-Rich Acidic Grasslands. Water Air and
  Soil Pollution 224.
- Duprè, C., C. J. Stevens, T. Ranke, A. Bleeker, C. Peppler-Lisbach, D. J. G. Gowing, N. B. Dise, E.
   Dorland, R. Bobbink, and M. Diekmann. 2010. Changes in species richness and composition in European acidic grasslands over the past 70 years: the contribution of cumulative atmospheric nitrogen deposition. Global Change Biology 16:344-357.

50 Ellenberg, H., H. E. Weber, R. Dull, V. Wirth, W. Werner, and D. Paulissen. 1991. Zeigerwerte von 51 Pflanzen in Mitteleuropa. Scripta Geobotanica **18**:1-248.

- Falkengren, G. U., J. Brunet, and M. Diekmann. 1998. Nitrogen mineralisation in deciduous forest soils in south Sweden in gradients of soil acidity and deposition. Environmental-Pollution. **102**:415-420.
- Falkengrengrerup, U. and H. Lakkenborgkristensen. 1994. Importance of ammonium and nitrate to the performance of herb-layer species from deciduous forests in Southern Sweden. Environmental and Experimental Botany **34**:31-38.
- Field, C., N. B. Dise, R. J. Payne, A. Britton, B. A. Emmett, R. Helliwell, S. Hughes, L. M. Jones, J. R. Leake, G. Phoenix, S. Power, L. Sheppard, G. Southon, C. Stevens, and S. J. M. Caporn. *in press*. Nitrogen drives plant community change across semi-natural habitats. Ecosystems.
- Field, C. D., N. B. Dise, R. J. Payne, A. J. Britton, B. A. Emmett, R. C. Helliwell, S. Hughes, L. Jones, S. Lees, J. R. Leake, I. D. Leith, G. K. Phoenix, S. A. Power, L. J. Sheppard, G. E. Southon, C. J. Stevens, and S. J. M. Caporn. 2014. The Role of Nitrogen Deposition in widespread Plant Community Change Across Semi-natural Habitats. Ecosystems.
- Friedrich, U., G. von Oheimb, C. Dziedek, W.-U. Kriebitzsch, K. Selbmann, and W. Haerdtle. 2011. Mechanisms of purple moor-grass (Molinia caerulea) encroachment in dry heathland ecosystems with chronic nitrogen inputs. Environmental Pollution **159**:3553-3559.
- Gujarati, D. N. 1995. Basic Econometrics. 3 edition. McGraw-Hill, New York.
- Henrys, P. A., C. J. Stevens, S. M. Smart, L. C. Maskell, K. J. Walker, C. D. Preston, A. Crowe, E. C. Rowe, D. J. Gowing, and B. A. Emmett. 2011. Impacts of nitrogen deposition on vascular plants in Britain: an analysis of two national observation networks. Biogeosciences **8**:3501-3518.
- Hill, M. O., C. D. Preston, and D. B. Roy. 2004. PLANTATT- Attributes of British and Irish Plants: Status, Size, Life History, Geography and Habitats. Biological Records Centre, NERC Centre for Ecology and Hydrology, Monkswood.
- Horswill, P., O. O'Sullivan, G. K. Phoenix, J. A. Lee, and J. R. Leake. 2008. Base cation depletion, eutrophication and acidification of species-rich grasslands in response to long-term simulated nitrogen deposition. Environmental Pollution **155**:336-349.
- Jones, L., A. Provins, M. Holland, G. Mills, F. Hayes, B. Emmett, J. Hall, L. Sheppard, R. Smith, M. Sutton, K. Hicks, M. Ashmore, R. Haines-Young, and L. Harper-Simmonds. 2014. A review and application of the evidence for nitrogen impacts on ecosystem services. Ecosystem Services 7:76-88
- Jones, M. L. M., H. L. Wallace, D. Norris, S. A. Brittain, S. Haria, R. E. Jones, P. M. Rhind, B. R. Reynolds, and B. A. Emmett. 2004. Changes in vegetation and soil characteristics in coastal sand dunes along a gradient of atmospheric nitrogen deposition. Plant Biology **6**:598-605.
- Kirk, G. J. D., P. H. Bellamy, and R. M. Lark. 2010. Changes in soil pH across England and Wales in response to decreased acid deposition. Global Change Biology **16**:3111-3119.
- 88 Maskell, L. C., S. M. Smart, J. M. Bullock, K. Thompson, and C. J. Stevens. 2010. Nitrogen deposition 89 causes widespread loss of species richness in British habitats. Global Change Biology **16**:671-90 679.
  - McClean, C. J., L. J. L. van den Berg, M. R. Ashmore, and C. D. Preston. 2011. Atmospheric nitrogen deposition explains patterns of plant species loss. Global Change Biology **17**:2882-2892.
    - McGovern, S., C. D. Evans, P. Dennis, C. Walmsley, and M. A. McDonald. 2011. Identifying drivers of species compositional change in a semi-natural upland grassland over a 40-year period. Journal of Vegetation Science **22**:346-356.
- Paulissen, M. P. C. P., P. J. M. van der Ven, A. J. Dees, and R. Bobbink. 2004. Differential effects of
   nitrate and ammonium on three fen bryophyte species in relation to pollutant nitrogen
   input. New Phytologist 164:451-458.

- Payne, R. J., N. B. Dise, C. J. Stevens, D. J. Gowing, and B. Partners. 2013. Impact of nitrogen
   deposition at the species level. Proceedings of the National Academy of Sciences of the
   United States of America 110:984-987.
- Provoost, S., M. L. M. Jones, and S. E. Edmondson. 2011. Changes in landscape and vegetation of coastal dunes in northwest Europe: a review. Journal of Coastal Conservation **15**:207-226.
- 104 R\_Development\_Core\_Team. 2008. R: a language and environment for statistical computing. R
  105 Foundation for Statistical Computing, Vienna, Austria.

106

107

108

114

115

116

117118

119

120

121

122

123

124

125

126

127

128129

130

131

132

133

134

135

- Remke, E., E. Brouwer, A. Kooijman, I. Blindow, and J. G. M. Roelofs. 2009. Low Atmospheric Nitrogen Loads Lead to Grass Encroachment in Coastal Dunes, but Only on Acid Soils. Ecosystems **12**:1173-1188.
- Roem, W. J. and F. Berendse. 2000. Soil acidity and nutrient supply ratio as possible factors determining changes in plant species diversity in grassland and heathland communities.

  Biological Conservation **92**:151-161.
- RoTAP. 2012. Review of Transboundary Air Pollution (RoTAP): Acidification, Eutrophication, Ground Level Ozone and Heavy Metals in the UK Centre for Ecology & Hydrology, Edinburgh.
  - Seed, L., P. Wolseley, L. Gosling, L. Davies, and S. A. Power. 2013. Modelling relationships between lichen bioindicators, air quality and climate on a national scale: Results from the UK OPAL air survey. Environmental Pollution **182**:437-447.
  - Sheppard, L. J., I. D. Leith, T. Mizunuma, J. N. Cape, A. Crossley, S. Leeson, M. A. Sutton, N. van Dijk, and D. Fowler. 2011. Dry deposition of ammonia gas drives species change faster than wet deposition of ammonium ions: evidence from a long-term field manipulation. Global Change Biology **17**:3589-3607.
  - Sheppard, L. J., I. D. Leith, T. Mizunuma, S. Leeson, S. Kivimaki, J. N. Cape, N. van Dijk, D. Leaver, M. A. Sutton, D. Fowler, L. J. L. Van den Berg, A. Crossley, C. Field, and S. Smart. 2014. Inertia in an ombrotrophic bog ecosystem in response to 9 years' realistic perturbation by wet deposition of nitrogen, separated by form. Global Change Biology **20**:566-580.
  - Smart, S. M., M. R. Ashmore, M. Hornung, W. A. Scott, D. A. Fowler, U. Dragosits, D. C. Howard, M. A. Sutton, and D. Famulari. 2004. Detecting the Signal of Atmospheric N Deposition in Recent National-Scale Vegetation Change Across Britain. Water, Air, & Soil Pollution: Focus **4**:269-278.
  - Smart, S. M., P. A. Henrys, B. V. Purse, J. M. Murphy, M. J. Bailey, and R. H. Marrs. 2012. Clarity or confusion? Problems in attributing large-scale ecological changes to anthropogenic drivers. Ecological Indicators **20**:51-56.
  - Smart, S. M. and W. A. Scott. 2004. Bias in Ellenberg indicator values problems with detection of the effect of vegetation type. Journal of Vegetation Science **15**:843-846.
  - Southon, G. E., C. Field, S. J. M. Caporn, A. J. Britton, and S. A. Power. 2013. Nitrogen Deposition Reduces Plant Diversity and Alters Ecosystem Functioning: Field-Scale Evidence from a Nationwide Survey of UK Heathlands. Plos One 8.
- Stevens, C. J., N. B. Dise, D. J. G. Gowing, and J. O. Mountford. 2006. Loss of forb diversity in relation to nitrogen deposition in the UK: regional trends and potential controls. Global Change Biology **12**:1823-1833.
- Stevens, C. J., N. B. Dise, J. O. Mountford, and D. J. Gowing. 2004. Impact of nitrogen deposition on the species richness of grasslands. Science **303**:1876-1879.
- Stevens, C. J., C. Dupre, E. Dorland, C. Gaudnik, D. J. G. Gowing, A. Bleeker, M. Diekmann, D. Alard, R.
   Bobbink, D. Fowler, E. Corcket, J. O. Mountford, V. Vandvik, P. A. Aarrestad, S. Muller, and N.
   B. Dise. 2010a. Nitrogen deposition threatens species richness of grasslands across Europe.
   Environmental Pollution 158:2940-2945.
- Stevens, C. J., P. Manning, L. J. L. van den Berg, M. C. C. de Graaf, G. W. W. Wamelink, A. W. Boxman,
   A. Bleeker, P. Vergeer, M. Arroniz-Crespo, J. Limpens, L. P. M. Lamers, R. Bobbink, and E.
   Dorland. 2011. Ecosystem responses to reduced and oxidised nitrogen inputs in European
   terrestrial habitats. Environmental Pollution 159:665-676.

150	Stevens, C. J., K. Thompson, J. P. Grime, C. J. Long, and D. J. G. Gowing. 2010b. Contribution of
151	acidification and eutrophication to declines in species richness of calcifuge grasslands along
152	a gradient of atmospheric nitrogen deposition. Functional Ecology 24:478-484.
153	van den Berg, L. J. L., E. Dorland, P. Vergeer, M. A. C. Hart, R. Bobbink, and J. G. M. Roelofs. 2005.
154	Decline of acid-sensitive plant species in heathland can be attributed to ammonium toxicity
155	in combination with low pH. New Phytologist <b>166</b> :551-564.
156	van den Berg, L. J. L., C. J. H. Peters, M. R. Ashmore, and J. G. M. Roelofs. 2008. Reduced nitrogen has
157	a greater effect than oxidised nitrogen on dry heathland vegetation. Environmental Pollution
158	<b>154</b> :359-369.
159	van den Berg, L. J. L., P. Vergeer, T. C. G. Rich, S. M. Smart, D. Guest, and M. R. Ashmore. 2011. Direct
160	and indirect effects of nitrogen deposition on species composition change in calcareous
161	grasslands. Global Change Biology 17:1871-1883.
162	Verheyen, K., L. Baeten, P. De Frenne, M. Bernhardt-Roemermann, J. Brunet, J. Cornelis, G. Decocq,
163	H. Dierschke, O. Eriksson, R. Hedl, T. Heinken, M. Hermy, P. Hommel, K. Kirby, T. Naaf, G.
164	Peterken, P. Petrik, J. Pfadenhauer, H. Van Calster, GR. Walther, M. Wulf, and G.
165	Verstraeten. 2012. Driving factors behind the eutrophication signal in understorey plant
166	communities of deciduous temperate forests. Journal of Ecology 100:352-365.
167	Vile, D., B. Shipley, and E. Garnier. 2006. Ecosystem productivity can be predicted from potential
168	relative growth rate and species abundance. Ecology Letters 9:1061-1067.
1.00	
169	
170	
171	

### 1 Figure legends

20

21

22

23

- 2 Figure 1 Effect sizes (slopes) of total N deposition on the number of species for UK habitats.
- 3 Only significant effects are shown. Actual ranges of N deposition covered by each vegetation
- 4 type differs but slopes are shown for a range between 5 and 45 kgNha<sup>-1</sup>y<sup>-1</sup>.
- 5 Figure 2 Effect sizes (slopes) of NH<sub>x</sub>:NO<sub>y</sub> ratio in deposition on the number of species for UK
- 6 habitats. Only significant effects are shown. Actual ranges of NH<sub>x</sub>:NO<sub>y</sub> ratio covered by each
- 7 vegetation type differs but slopes are shown for a range between 0.5 and 3.5.
- 8 Figure 3 Effect sizes (slopes) of total N deposition on the Ellenberg N number for UK
- 9 habitats. Only significant effects are shown. Actual ranges of N deposition covered by each
- vegetation type differs but slopes are shown for a range between 5 and 45 kgNha<sup>-1</sup>y<sup>-1</sup>.
- 11 Figure 4 Effect sizes (slopes) of NH<sub>x</sub>:NO<sub>y</sub> ratio in deposition on the Ellenberg N number for
- 12 UK habitats. Only significant effects are shown. Actual ranges of NH<sub>x</sub>:NO<sub>y</sub> ratio covered by
- each vegetation type differs but slopes are shown for a range between 0.5 and 3.5.
- 14 Figure 5 Effect sizes (slopes) of total N deposition on the Grass:Forb ratio for UK habitats.
- Only significant effects are shown. Actual ranges of N deposition covered by each vegetation
- type differs but slopes are shown for a range between 5 and 45 kgNha<sup>-1</sup>y<sup>-1</sup>.
- 17 Figure 6 Effect sizes (slopes) of NH<sub>x</sub>:NO<sub>y</sub> ratio in deposition on the Grass:Forb ratio for UK
- habitats. Only significant effects are shown. Actual ranges of NH<sub>x</sub>:NO<sub>y</sub> ratio covered by each
- vegetation type differs but slopes are shown for a range between 0.5 and 3.5.

- 1 Table 1 Coefficients and their significance for the habitats dry upland heaths, dry lowland heaths,
- 2 bogs & mires and base rich mires. Results of a multiple regression (A) and regression analysis of the
- 3 residuals of models containing either NH<sub>x</sub> or NO<sub>y</sub> with respectively NO<sub>y</sub> and NH<sub>x</sub> to separate N form
- 4 (B) (see methods for details). Results are shown for the response variables: species richness,
- 5 Ellenberg N fertility score and grass:forb ratio. Grass:forb ratio was log transformed.

			Sp. rich	Ellenberg N	Grass:Forb
Dry upland heaths	Α	Temperature	-0.002	0.017	0.113
		Precipitation	0.014***	-0.001**	0.000
		S deposition	0.103	-0.017	-0.003
		Total N deposition	-0.174***	0.006	0.031*
		NHx:NOy ratio	0.783	0.475***	0.528
	В	NOy deposition	-0.037	-0.007	-0.001
		NHx deposition	-0.019	0.009	0.012
Dry lowland heaths	Α	Temperature	0.474	0.103**	0.035
		Precipitation	0.010	-0.002**	0.005*
		S deposition <sup>1</sup>	-0.320	-0.480	1.290
		Total N deposition	-0.266***	-0.022***	0.035*
		NHx:NOy ratio	1.148	0.634***	0.342
	В	NOy deposition	-0.099	-0.021*	0.001
		NHx deposition	-0.030	0.006	0.011
Bogs and Mires	Α	Temperature	-0.226*	0.032*	0.178***
		Precipitation	0.011***	0.000	0.002**
		S deposition	-0.033	0.001	-0.030*
		Total N deposition	-0.057***	0.007**	0.028***
		NHx:NOy ratio	0.797	0.374***	0.312
	В	NOy deposition	-0.044	-0.006	0.000
		NHx deposition	0.004	0.007*	0.009
Base rich Mires	Α	Temperature	-0.444	0.125**	0.107**
		Precipitation	0.001	-0.004***	-0.003***
		S deposition	0.004	0.023*	0.016
		Total N deposition	-0.011	0.016*	0.005
		NHx:NOy ratio	-0.364	-0.046	0.081
	В	NOy deposition	0.004	0.004	0.003
		NHx deposition	-0.014	0.008	-0.001

Number of plots: Dry upland heaths (267), dry lowland heaths (182), Bogs and Mires (1136), Base rich Mires (274). <sup>1</sup> S deposition was

<sup>7</sup> inverse transformed

- 8 Table 2 Coefficients and their significance for the habitats acidic grasslands, mesotrophic grasslands,
- 9 calcareous grassland and woodlands. Results of a multiple regression (A) and regression analysis of
- the residuals of models containing either NH<sub>x</sub> or NO<sub>y</sub> with respectively NO<sub>y</sub> and NH<sub>x</sub> to separate N
- form (B) (see methods for details). Results are shown for the response variables: species richness,
- 12 Ellenberg N fertility score and grass:forb ratio. Grass:forb ratio was log transformed.

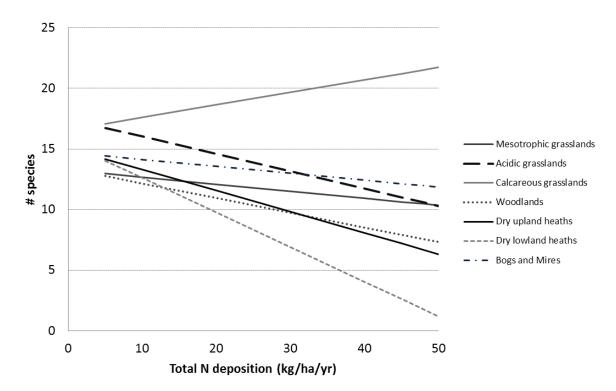
			Sp. rich	Ellenberg N	Grass:Forb
Acidic grasslands	Α	Temperature	0.097	0.086***	-0.098**
		Precipitation <sup>1</sup>	-0.003***	0.016***	-0.040
		S deposition	-0.113**	0.005	-0.013
		Total N deposition	-0.145***	-0.015***	0.030***
		NHx:NOy ratio	-1.783***	0.368***	0.455**
	В	NOy deposition	-0.011	-0.018***	0.002
		NHx deposition	-0.061*	0.006	-0.001
Mesotrophic grasslands	Α	Temperature	0.457***	0.010	-0.110***
		Precipitation	0.028***	-0.006***	0.000
		S deposition	0.006	0.008*	0.000
		Total N deposition	-0.058*	0.008**	-0.001
		NHx:NOy ratio	-0.947**	0.172***	0.100
	В	NOy deposition	0.023	-0.010	-0.001
		NHx deposition	-0.096***	0.018***	0.003
Calcareous grasslands	Α	Temperature	-0.906***	0.126***	0.015
		Precipitation	-0.012***	-0.005***	0.004***
		S deposition	-0.034	0.018*	0.031*
		Total N deposition	0.103**	0.032***	-0.027***
		NHx:NOy ratio	-2.849***	0.618***	0.258*
	В	NOy deposition	0.259***	-0.025**	-0.026
		NHx deposition	-0.109**	0.046***	-0.002
Woodlands	Α	Temperature	-0.103	0.207***	-0.145*
		Precipitation	0.002	-0.006***	0.003
		S deposition	-0.044	0.008	-0.007
		Total N deposition	-0.121***	0.006	-0.003
		NHx:NOy ratio	1.178*	0.271**	-0.116
	В	NOy deposition	-0.229***	-0.022	0.017
		NHx deposition	0.019	0.026***	-0.015

Number of plots: Acidic grasslands (1090), Mesotrophic grasslands (1195), Calcareous grasslands (830), Woodlands (514). <sup>1</sup> Precipitation

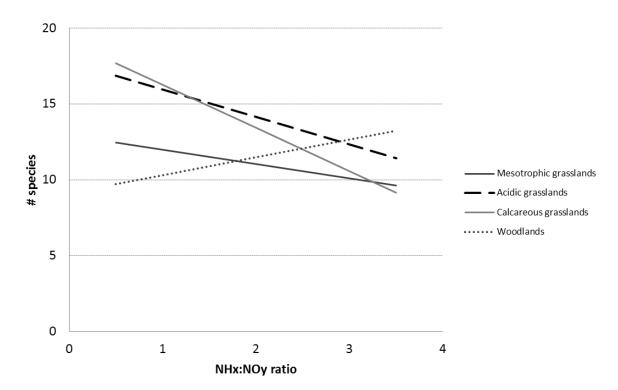
14 was inverse transformed

13

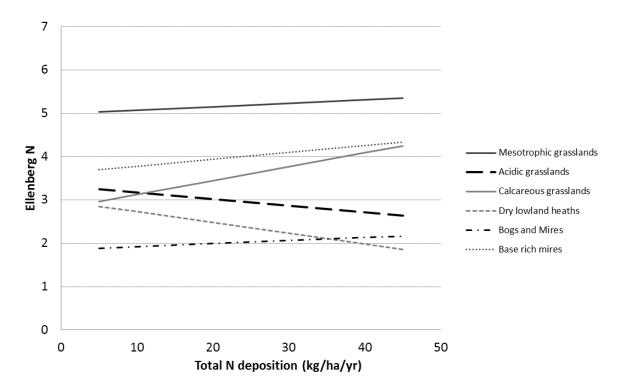
## 1 Figure 1



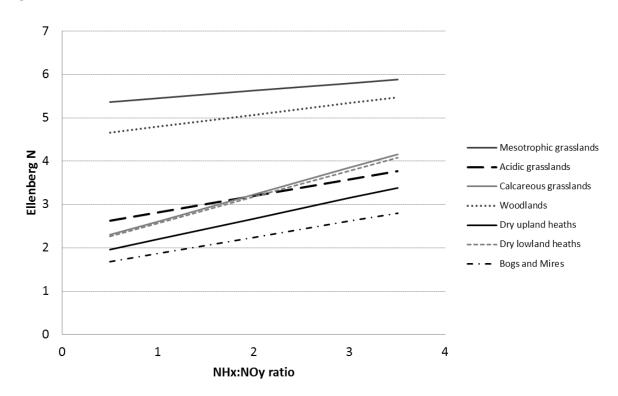
## 3 Figure 2



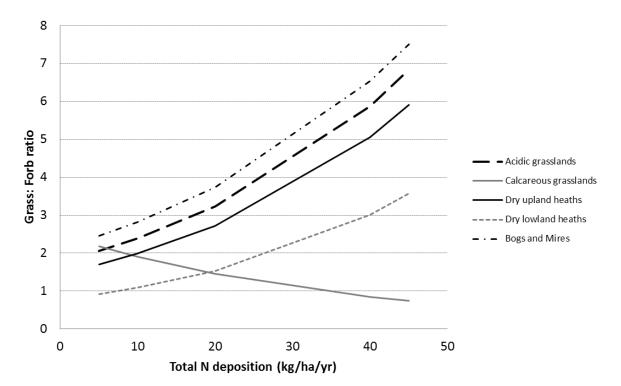
## 6 Figure 3



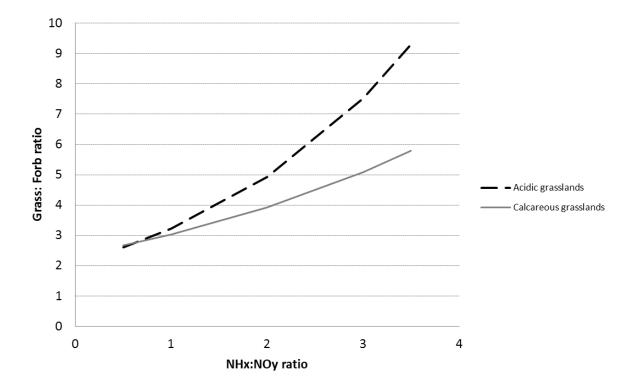
### 8 Figure 4



## 11 Figure 5



13 Figure 6



- 1 This is an author-created version. The full article can be found at: Environmental Pollution
- 2 October 2015:DOI: 10.1016/j.envpol.2015.09.017
- 3 **Supplementary material**
- 4 Table 1: number of plots included in the analysis

NVC type	# plots included in the analysis
Dry upland Heaths	267
Dry lowland Heaths	182
Bogs and Mires	1136
Base rich Mires	274
Acidic grasslands	1090
Mesotrophic grasslands	1195
Calcareous grasslands	869
Woodlands	514