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4 **Evidence for differential effects of reduced and oxidised nitrogen deposition on**
5 **vegetation independent of nitrogen load**

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22 Capsule:

23 Effects of total N deposition and reduced and oxidised N deposition were studied across
24 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.

31

32 **Abstract**

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
38 national monitoring data in the UK to provide an extensive evaluation of whether there are
39 differences in the effects of reduced and oxidised N deposition across eight habitat types
40 (acid, calcareous and mesotrophic grasslands, upland and lowland heaths, bogs and mires,
41 base-rich mires, woodlands). We analysed data from 6860 plots in the British Countryside
42 Survey 2007 for effects of total N deposition and N form on species richness, Ellenberg N
43 values and grass:forb ratio. Our results provide clear evidence that that N deposition affects
44 species richness in all habitats except base-rich mires, after factoring out correlated
45 explanatory variables (climate and sulphur deposition). In addition, the form of N in
46 deposition appears important for the biodiversity of grasslands and woodlands but not
47 mires and heaths. Ellenberg N increased more in relation to NH_x deposition than NO_y
48 deposition in all but one habitat type. Relationships between species richness and N form
49 were habitat-specific: acid and mesotrophic grasslands appear more sensitive to NH_x
50 deposition while calcareous grasslands and woodlands appeared more responsive to NO_y
51 deposition. These relationships are likely driven by the preferences of the component plant
52 species for oxidised or reduced forms of N, rather than by soil acidification.

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59 **Keywords**

60 $\text{NH}_x:\text{NO}_y$ ratio, N deposition, countryside survey, acidification, grassland, heathland, bogs

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1 Introduction

2

3 There is widespread evidence across the globe, from both experiments and field surveys, of
4 the significant ecological impacts of nitrogen (N) deposition on semi-natural ecosystems of
5 low nutrient status (e.g. Bobbink et al. 2010), which also carries economic costs (Jones et al.
6 2014). However, interpretation and quantification of these effects, and predictions of the
7 benefits of emission control policies, need to consider the different components of N
8 deposition (Brink et al. 2011). There are two main chemical forms – reduced N (ammonia,
9 NH_3 and ammonium, NH_4^+) emitted primarily from agricultural sources, and oxidised N
10 (nitrogen oxides, NO_x , nitric acid, HNO_3^- and nitrate NO_3^-) emitted primarily from fossil fuel
11 combustion. In addition, N deposition may be in the form of dry deposition of gases and
12 aerosols, which is most important close to sources, and in regions of the world with low
13 rainfall, and as wet deposition as snow, dew, cloud or rainwater, which are important in
14 more remote regions and in areas with high rainfall.

15 The mechanisms underlying the ecological effects of N deposition include direct toxicity,
16 growth stimulation and competitive exclusion, soil acidification and increased susceptibility
17 to other abiotic and biotic stresses (e.g. Bobbink et al. 1998, Roem and Berendse 2000).
18 There are strong reasons, which have been recently reviewed by (Stevens et al. 2011), for
19 expecting that there may be different effects of reduced and oxidised N deposition for each
20 of these mechanisms. For example, foliar uptake of gaseous NH_3 is more likely to be directly
21 toxic than uptake of gaseous nitrogen oxides, while soil NH_4^+ is more likely to be toxic to
22 plant roots than soil NO_3^- (Sheppard et al. 2011, Sheppard et al. 2014). Plant species also
23 differ strongly in their preference and tolerance for NH_4^+ or NO_3^- uptake from soil solution
24 with species of acidic habitats generally more tolerant of higher soil ammonium
25 (Falkengrengrerup and Lakkenborgkristensen 1994). The soil $\text{NH}_4^+/\text{NO}_3^-$ ratio is partly a
26 function of the ratio in atmospheric deposition, but also of the degree of nitrification in
27 soils; high rates of nitrification result in a lower soil solution $\text{NH}_4^+/\text{NO}_3^-$ ratio, which may
28 reduce the risk of direct NH_4^+ toxicity but may increase acidification because of the greater
29 oxidation to NO_3^- .

30 Experimental studies provide some evidence of the differential effects of reduced and
31 oxidised N deposition. For example, van den Berg et al. (2008) showed that higher
32 $\text{NH}_4^+/\text{NO}_3^-$ ratios in deposition to heathland mesocosms had significant adverse effects on
33 acid-sensitive species but not on acid-tolerant species that were also tolerant of high soil
34 $\text{NH}_4^+/\text{NO}_3^-$ ratios. This effect was lost in limed mesocosms, suggesting that acidification at
35 higher $\text{NH}_4^+/\text{NO}_3^-$ ratios was the key driving mechanism. By contrast, in Mediterranean
36 maquis vegetation, the application of both NH_4^+ and NO_3^- increased biomass but not plant
37 diversity, while NH_4^+ alone increased plant diversity but not biomass (Dias et al. 2014); these
38 effects can at least partly be explained by the different responses of individual species to
39 total N inputs or to reduced N deposition specifically.

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

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

59 These and other findings from field surveys suggest that the responses to N deposition of
60 vegetation characteristics in different habitats may be at least partly explained by
61 differences in the underlying mechanisms of impact of reduced and oxidised N, mediated by
62 soil pH, with acidification effects prevailing in poorly-buffered habitats and eutrophication
63 effects in well-buffered habitats. Few field surveys have tried to separately evaluate the
64 strength of associations with reduced and oxidised nitrogen but were only able to do so
65 with relatively low number of samples/sites (Caporn et al. 2014, Field et al. 2014). Three
66 studies have showed adverse changes in vegetation composition that were significantly
67 correlated with reduced N deposition but not with oxidised N deposition: an increase in
68 mean Ellenberg fertility index in semi-natural grassland and heaths/bogs between 1990 and
69 1998 in UK Countryside Survey data (Smart et al. 2004); a loss of species with a low
70 Ellenberg fertility index in UK national recording data between 1987 and 1999 (McClellan et
71 al. 2011); and increases in graminoid cover and decreases in lichen cover in heathlands
72 (Southon et al. 2013). A further study showed effects only of dry deposition of NH_x and no
73 effect of wet reduced or oxidised N on abundance of N sensitive epiphytic lichens (Seed et
74 al. 2013).

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

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

91 -The form of N (oxidised NO_y , or reduced NH_x) in deposition has an effect on vegetation
92 composition that is independent of, and additional to, that of total N deposition.

93 -Reduced N deposition has a greater impact on vegetation composition than oxidised N
94 deposition.

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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
10 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
13 supplementary material). Earlier studies have shown that soil pH or base saturation can
14 explain species richness and can affect the responses of the vegetation and the ecosystem
15 to N deposition (van den Berg et al. 2005, Stevens et al. 2006). Therefore, habitats that
16 belong to a similar broad NVC classification but differ strongly in average soil pH and/or
17 base cation content (heaths, mires and grasslands) were subdivided according to pH for
18 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
22 fen habitat consist of the NVC classes M22 to M38. Upland dry heaths are NVC classes H10
23 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
25 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
28 are known to affect species richness (Cleland et al. 2013). Average annual temperature (°C)
29 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
32 may thereby affect species richness (McGovern et al. 2011). Historical data on S deposition
33 was therefore included in our models to account for potential legacy of soil acidification
34 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
37 deposition ranged from 5.1 to 54.2 kg N ha⁻¹yr⁻¹ while S deposition ranged from 5.0 to 43.5
38 kg S ha⁻¹yr⁻¹. Oxidised and reduced N were included in our models as the sum of wet and dry

39 NO_y or NH_x deposition and expressed in kg N ha⁻¹yr⁻¹. NO_y deposition ranged from 2.5 to
40 25.6 kg N ha⁻¹yr⁻¹, NH_x deposition ranged from 2.3 to 36.1 kg N ha⁻¹yr⁻¹. The ranges of N and
41 S deposition for each habitat are different and depend on their geographical distribution.

42 *Linear models*

43 Multicollinearity is common between variables such as N deposition and the different forms
44 of N in deposition, S deposition and climatic variables. In our dataset, NH_x and NO_y
45 deposition were highly correlated ($r=0.69$, $p<0.001$) and could therefore not be analysed
46 simultaneously. In addition, total N deposition and either NH_x or NO_y 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 multicollinearity and that
50 each test specific hypotheses. Linearity of the relationship between the predictor factors
51 and the dependent variables were tested in single linear regressions. If needed, data were
52 transformed to meet assumptions of linearity.

53 In the first method (models coded with A), the effect of NH_x/NO_y ratio in deposition was
54 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
57 variables: total N deposition, NH_x/NO_y ratio, S deposition, precipitation and temperature. All
58 models were at first explicitly tested for spatial autocorrelation in the response variable and
59 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
61 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
63 distances (using Pythagoras theorem) between sites with coordinates given by Easting and
64 Northing. When spatial autocorrelation was not present or not severe, multiple linear
65 regression models were used.

66 In the second method (models coded with B), the additional effects of either NH_x or of NO_y
67 on species richness, Ellenberg N and grass:forb ratio were tested after taking into account
68 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_x, S deposition,
70 precipitation and temperature were regressed in a second model against NO_y. i.e. the
71 relationship between NO_y deposition and the unexplained variance of the model was tested.
72 The calculation was then repeated for an analysis of NH_x on the residuals of a model that
73 included NO_y. Given that the data cover a substantial range of NO_y and NH_x deposition, and
74 making the assumption that the observed responses with N deposition are linear, any
75 differential effect of reduced and oxidised N deposition is independent of the range of the
76 length of the deposition gradient. In this way, the slope coefficients (effect sizes) that are

77 derived allow a comparison of the independent effects of either NH_x and NO_y, after
78 accounting for other sources of (co-correlated) variation.

79

80 Multicollinearity in the models was detected by calculating the Pearson correlation
81 coefficient among pairs of the predictors and by calculating the variance inflation factors
82 (VIF) for each predictor in the model. Predictors with VIF of less than 4 were maintained in
83 the models since these indicate that problems with multicollinearity are not severe (Gujarati
84 1995). Predictor variables that were highly correlated (VIF>4) were not analysed in the same
85 model. In an additional step, multicollinearity was explored by comparing the beta-
86 coefficients of the explanatory variables that were obtained in a multiple regression with
87 the beta coefficients from single regressions of these explanatory variables. In this analysis,
88 major changes in beta coefficient or changes in sign indicate multicollinearity between
89 explanatory variables that needs to be accounted for.

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

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_x/NO_y 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_x/NO_y ratio was
11 found, while there was no significant effect of N form on the upland and lowland heathlands
12 and the base-rich mires, bogs and mires. Analyses of residuals against NH_x and NO_y (Models
13 B) in all cases were consistent with responses shown by NH_x/NO_y ratio (Models A). These
14 analyses showed that the negative effects on species richness in all three grassland habitats
15 were driven by strong negative effects of NH_x. Species richness in calcareous grasslands was
16 also positively related to NO_y (Table 2). In woodlands in contrast, the negative effects on
17 species richness were strongly related to NO_y deposition.

18 Total N deposition increased Ellenberg fertility index for bogs & mires, base rich mires,
19 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_x/NO_y
23 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_y 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_x
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
31 & mires and acidic grasslands (Figure 5). Only in calcareous grasslands, grass:forb ratio was
32 found to be lower with increased N deposition. There was no effect on grass:forb ratio in
33 base rich mires, mesotrophic grasslands or woodlands. In acidic and calcareous grasslands
34 the increased grass:forb ratio was associated with increased NH_x/NO_y ratio (Table 2, Figure
35 6), but separate relationships for either reduced N or oxidised N were not significant and
36 could not be used to infer which form of N was more responsible.

37 *Additional environmental factors*

38 Climate variables were frequently a significant explanatory variable for total species richness
39 and Ellenberg fertility score (Table 1 and 2). Precipitation was negatively associated with
40 species richness in the acidic and calcareous grasslands but positively associated with
41 species richness in mesotrophic grasslands, dry upland heath and bogs & mires. A higher
42 precipitation was associated with a lower Ellenberg fertility score in most habitats.
43 Temperature was positively associated with Ellenberg fertility scores. Past sulphur
44 deposition also showed some significant effects. Sulphur deposition showed a significant
45 relationship with species richness in acidic grasslands (negative), Ellenberg fertility scores for
46 mesotrophic and calcareous grasslands (positive) and grass:forb ratio for bogs & mires
47 (negative). This highlights the importance of factoring out these co-correlated variables to
48 genuinely extract any relationships due to N deposition or N form.

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1 Discussion

2 *Total N deposition effects on species richness*

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

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

31

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

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

47

48 Base rich habitats and, to a lesser extent, bogs and mires increased in Ellenberg fertility
49 index with increasing N deposition which was not necessarily accompanied with a loss in
50 species richness suggesting that elevated N deposition in these habitats results in a shift in
51 species composition favouring more nutrient-loving species. In contrast, acid grasslands and
52 lowland heaths show a small decline in fertility index with increasing N deposition. Others
53 have also reported lower Ellenberg N values with higher N deposition in acid grasslands
54 (Maskell et al. 2010) and heathlands (Caporn et al. 2014) and these relationships may be
55 linked to the high correlation between Ellenberg N and Ellenberg R, suggesting mechanisms
56 such as acidification to operate in these systems. The exact mechanisms for these
57 relationships are however not known and need further exploration at the species level of
58 both vascular plants and bryophytes; studies have shown much greater effects of N
59 deposition on bryophyte species richness than vascular plant species richness (e.g. Caporn
60 et al. 2014).

61 *N form and the relative influence of reduced versus oxidised N*

62 NH_x deposition and NO_y deposition are highly correlated. In addition, NH_x was highly
63 correlated to total N deposition. Separate analysis of NH_x and NO_y effects in models that
64 allowed us to factor out the variance that was explained by either one of the N forms was
65 therefore considered the best method to compare effects of these N forms, after taking
66 account of other variables and the multicollinearity that existed in the datasets. Since both
67 forms are correlated the variance that is explained by one N form, and which is factored out,
68 is likely to contain some degree of variance that in fact should be attributed to the other
69 form. The method that we employed here is therefore considered conservative in its
70 estimation of effect sizes and significance levels.

71 The range of NH_x (2.3 - 36.1 $\text{kgNha}^{-1}\text{y}^{-1}$) exceeds that of NO_y (2.5 - 25.6 $\text{kgNha}^{-1}\text{y}^{-1}$) over all
72 habitats together. Although the gradient length of explanatory variables may affect the
73 outcome of the analysis in small data sets (Smart and Scott 2004), large datasets such as the
74 CS data capture a good proportion of the relationship (i.e. not just a small segment), even
75 with smaller ranges of the explanatory variable. Our analysis is based on the assumption
76 that the relationships are linear (transformations were applied when necessary) between
77 the response variable and either NH_x or NO_y and estimations of the effect sizes of the
78 relationships are therefore considered relatively unaffected by the length of the gradients.
79 Plots of beta coefficients of the regressions for NO_y and NH_x against N-gradient length

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

86
87 The form of N in deposition independent of N load did affect species richness, but only in
88 certain habitats. N form altered species richness in all grasslands and in woodlands but not
89 in the mires and heaths. The lack of response of species richness and also grass:forb ratio, to
90 N form in heaths and mires may be due to the prevailing acidic conditions, restraining
91 nitrification rates with naturally high soil $\text{NH}_4^+/\text{NO}_3^-$ ratios and low base cation
92 concentrations (e.g. De Graaf et al. 2009). Many species of acidic habitats, such as ericoids,
93 are generally adapted to elevated NH_4^+ concentrations and tolerate high NH_4^+
94 concentrations (De Graaf et al. 1998, Britto and Kronzucker 2002, Sheppard et al. 2014).

95
96 Where N form was important, in the acidic and mesotrophic grasslands NH_x appeared to be
97 more important than NO_y as a driver of species richness decline, corroborating an
98 experimental study on N form in acid grasslands (Dorland et al. 2013). However, in the
99 woodlands and in calcareous grassland, NO_x was more important than NH_y , having a positive
100 effect on species richness in the grassland but a negative effect in the woodland.
101 Nitrification and mineralisation in woodlands can be very high (Falkengrengrerup and
102 Lakkenborgkristensen 1994, Falkengren et al. 1998). Atmospheric deposition of oxidised N
103 may therefore favour nitrophilous species such as bramble and nettle that outcompete
104 slow-growing forb and shrub species that are more adapted to ammonium nutrition (such as
105 *Vaccinium myrtillus*), corroborating a recent simulation study (Stevens et al. this volume).
106 The positive impact of NO_y on calcareous grassland species richness may relate to the
107 preference of many calcareous species for available N in oxidised rather than in reduced
108 form.

109
110 The question remains why the species richness and composition of only some habitats are
111 sensitive to N form, even though the fertility index of almost all habitats responded to N
112 form. N form may alter species composition through preferences of the component species
113 for oxidised or reduced N, through direct toxicity of NH_3 and NH_4^+ (Britto and Kronzucker
114 2002, van den Berg et al. 2005, Sheppard et al. 2011), or through indirect effects mediated
115 by N-induced acidification (e.g. Bobbink et al. 1998, Stevens et al. 2011), which would be
116 more apparent in acidic habitats. The lack of significance of N form in the more acidic
117 habitats suggests that acidification is not the main cause. However, experimental studies
118 have shown that elevated $\text{NH}_4^+/\text{NO}_3^-$ ratios in deposition result in a decline of acid-sensitive
119 species but not of acid-loving species tolerant of high soil $\text{NH}_4^+/\text{NO}_3^-$ ratios (Paulissen et al.

120 2004, van den Berg et al. 2008). The response in acidic and mesotrophic grasslands may
121 therefore reflect the abundance of species that are sensitive to reduced N in these neutral
122 to moderately acidic habitats compared with the more strongly acidophile vegetation in
123 heaths, bogs and mires. Clearly responses to N form are habitat-specific, and may be driven
124 by the preference or tolerance of the component species for N in oxidised or reduced forms.

125

126 In conclusion, this study has shown that N affects species richness in almost all habitats,
127 after correlating factors such as temperature, rainfall and historical sulphur deposition have
128 been factored out. The form of N is important, with fertility index increasing with NH_x/NO_y
129 ratio in almost all habitats. However, the effects of the ratio on species richness were only
130 found in certain habitats (grasslands and woodland), not in others (mires and heaths). In
131 habitats where there were differential effects of one N form or the other, acidic and
132 mesotrophic grassland were more sensitive to NH_y , while calcareous grassland and
133 woodland were more sensitive to NO_x . This study suggests that, contrary to our original
134 hypothesis, sensitivity to N form is more likely due to the inherent preferences of
135 component species for oxidised or reduced N, rather than linked to soil acidification.
136 However, those preferences are related to soil pH with NH_4 -loving species generally more
137 prevalent on acidic soils.

138

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141 deposition data available.

142

143

144

1 References

- 2 Bennie, J., M. O. Hill, R. Baxter, and B. Huntley. 2006. Influence of slope and aspect on long-term
3 vegetation change in British chalk grasslands. *Journal of Ecology* **94**:355-368.
- 4 Bobbink, R., K. Hicks, J. Galloway, T. Spranger, R. Alkemade, M. Ashmore, M. Bustamante, S.
5 Cinderby, E. Davidson, F. Dentener, B. Emmett, J. W. Erisman, M. Fenn, F. Gilliam, A. Nordin,
6 L. Pardo, and W. De Vries. 2010. Global assessment of nitrogen deposition effects on
7 terrestrial plant diversity: a synthesis. *Ecological Applications* **20**:30-59.
- 8 Bobbink, R., M. Hornung, and J. G. M. Roelofs. 1998. The effects of air-borne nitrogen pollutants on
9 species diversity in natural and semi-natural European vegetation. *Journal of Ecology*
10 **86**:717-738.
- 11 Brink, C., H. van Grinsven, B. H. Jacobsen, A. Rabl, I.-M. Gren, M. Holland, Z. Klimont, K. Hicks, R.
12 Brouwer, R. Dickens, J. Willems, M. Termansen, G. Velthof, R. Alkemade, M. van Oorschot,
13 and J. Webb. 2011. Costs and benefits of nitrogen in the environment. Pages 513-540 in M.
14 A. Sutton, C. M. Howard, J. W. Erisman, G. Billen, A. Bleeker, P. Grennfelt, H. van Grinsven,
15 and B. Grizzetti, editors. *The European nitrogen assessment*. Cambridge University Press,
16 Cambridge.
- 17 Britto, D. T. and H. J. Kronzucker. 2002. NH₄⁺ toxicity in higher plants: a critical review. *Journal of*
18 *Plant Physiology* **159**:567-584.
- 19 Caporn, S. J. M., J. A. Carroll, N. B. Dise, and R. J. Payne. 2014. Impacts and indicators of nitrogen
20 deposition in moorlands: Results from a national pollution gradient study. *Ecological*
21 *Indicators* **45**:227-234.
- 22 Carey, P. D., S. Wallis, P. M. Chamberlain, A. Cooper, B. A. Emmett, L. C. Maskell, T. McCann, J.
23 Murphy, L. R. Norton, B. Reynolds, W. A. Scott, I. C. Simpson, S. M. Smart, and J. M. Ulllyett.
24 2008. *Countryside Survey: UK Results from 2007*. NERC/Centre for Ecology & Hydrology.
- 25 Cleland, E. E., S. L. Collins, T. L. Dickson, E. C. Farrer, K. L. Gross, L. A. Gherardi, L. M. Hallett, R. J.
26 Hobbs, J. S. Hsu, L. Turnbull, and K. N. Suding. 2013. Sensitivity of grassland plant community
27 composition to spatial vs. temporal variation in precipitation. *Ecology* **94**:1687-1696.
- 28 De Graaf, M. C. C., R. Bobbink, J. G. M. Roelofs, and P. J. M. Verbeek. 1998. Differential effects of
29 ammonium and nitrate on three heathland species. *Plant Ecology* **135**:185-196.
- 30 De Graaf, M. C. C., R. Bobbink, N. A. C. Smits, R. Van Diggelen, and J. G. M. Roelofs. 2009.
31 Biodiversity, vegetation gradients and key biogeochemical processes in the heathland
32 landscape. *Biological Conservation* **142**:2191-2201.
- 33 De Graaf, M. C. C., R. Bobbink, P. J. M. Verbeek, and J. G. M. Roelofs. 1997. Aluminium toxicity and
34 tolerance in three heathland species. *Water Air and Soil Pollution* **98**:229-239.
- 35 Dias, T., A. Clemente, M. A. Martins-Loucao, L. Sheppard, R. Bobbink, and C. Cruz. 2014. Ammonium
36 as a Driving Force of Plant Diversity and Ecosystem Functioning: Observations Based on 5
37 Years' Manipulation of N Dose and Form in a Mediterranean Ecosystem. *Plos One* **9**.
- 38 Diekmann, M. and U. Falkengren-Grerup. 2002. Prediction of species response to atmospheric
39 nitrogen deposition by means of ecological measures and life history traits. *Journal of*
40 *Ecology* **90**:108-120.
- 41 Dorland, E., C. J. Stevens, C. Gaudnik, E. Corcket, S. Rotthier, K. Wotherspoon, M. Jokerud, V.
42 Vandvik, M. B. Soons, M. M. Hefting, P. A. Aarrestad, D. Alard, M. Diekmann, C. Dupre, N. B.
43 Dise, D. J. G. Gowing, and R. Bobbink. 2013. Differential Effects of Oxidised and Reduced
44 Nitrogen on Vegetation and Soil Chemistry of Species-Rich Acidic Grasslands. *Water Air and*
45 *Soil Pollution* **224**.
- 46 Duprè, C., C. J. Stevens, T. Ranke, A. Bleeker, C. Peppeler-Lisbach, D. J. G. Gowing, N. B. Dise, E.
47 Dorland, R. Bobbink, and M. Diekmann. 2010. Changes in species richness and composition
48 in European acidic grasslands over the past 70 years: the contribution of cumulative
49 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.

52 Falkengren, G. U., J. Brunet, and M. Diekmann. 1998. Nitrogen mineralisation in deciduous forest
53 soils in south Sweden in gradients of soil acidity and deposition. *Environmental-Pollution*.
54 **102**:415-420.

55 Falkengrengrerup, U. and H. Lakkenborgkristensen. 1994. Importance of ammonium and nitrate to
56 the performance of herb-layer species from deciduous forests in Southern Sweden.
57 *Environmental and Experimental Botany* **34**:31-38.

58 Field, C., N. B. Dise, R. J. Payne, A. Britton, B. A. Emmett, R. Helliwell, S. Hughes, L. M. Jones, J. R.
59 Leake, G. Phoenix, S. Power, L. Sheppard, G. Southon, C. Stevens, and S. J. M. Caporn. *in*
60 *press*. Nitrogen drives plant community change across semi-natural habitats. *Ecosystems*.

61 Field, C. D., N. B. Dise, R. J. Payne, A. J. Britton, B. A. Emmett, R. C. Helliwell, S. Hughes, L. Jones, S.
62 Lees, J. R. Leake, I. D. Leith, G. K. Phoenix, S. A. Power, L. J. Sheppard, G. E. Southon, C. J.
63 Stevens, and S. J. M. Caporn. 2014. The Role of Nitrogen Deposition in widespread Plant
64 Community Change Across Semi-natural Habitats. *Ecosystems*.

65 Friedrich, U., G. von Oheimb, C. Dzedek, W.-U. Kriebitzsch, K. Selbmann, and W. Haerdle. 2011.
66 Mechanisms of purple moor-grass (*Molinia caerulea*) encroachment in dry heathland
67 ecosystems with chronic nitrogen inputs. *Environmental Pollution* **159**:3553-3559.

68 Gujarati, D. N. 1995. *Basic Econometrics*. 3 edition. McGraw-Hill, New York.

69 Henrys, P. A., C. J. Stevens, S. M. Smart, L. C. Maskell, K. J. Walker, C. D. Preston, A. Crowe, E. C.
70 Rowe, D. J. Gowing, and B. A. Emmett. 2011. Impacts of nitrogen deposition on vascular
71 plants in Britain: an analysis of two national observation networks. *Biogeosciences* **8**:3501-
72 3518.

73 Hill, M. O., C. D. Preston, and D. B. Roy. 2004. *PLANTATT- Attributes of British and Irish Plants:*
74 *Status, Size, Life History, Geography and Habitats*. Biological Records Centre, NERC Centre
75 for Ecology and Hydrology, Monkswood.

76 Horswill, P., O. O'Sullivan, G. K. Phoenix, J. A. Lee, and J. R. Leake. 2008. Base cation depletion,
77 eutrophication and acidification of species-rich grasslands in response to long-term
78 simulated nitrogen deposition. *Environmental Pollution* **155**:336-349.

79 Jones, L., A. Provins, M. Holland, G. Mills, F. Hayes, B. Emmett, J. Hall, L. Sheppard, R. Smith, M.
80 Sutton, K. Hicks, M. Ashmore, R. Haines-Young, and L. Harper-Simmonds. 2014. A review and
81 application of the evidence for nitrogen impacts on ecosystem services. *Ecosystem Services*
82 **7**:76-88.

83 Jones, M. L. M., H. L. Wallace, D. Norris, S. A. Brittain, S. Haria, R. E. Jones, P. M. Rhind, B. R.
84 Reynolds, and B. A. Emmett. 2004. Changes in vegetation and soil characteristics in coastal
85 sand dunes along a gradient of atmospheric nitrogen deposition. *Plant Biology* **6**:598-605.

86 Kirk, G. J. D., P. H. Bellamy, and R. M. Lark. 2010. Changes in soil pH across England and Wales in
87 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.

91 McClean, C. J., L. J. L. van den Berg, M. R. Ashmore, and C. D. Preston. 2011. Atmospheric nitrogen
92 deposition explains patterns of plant species loss. *Global Change Biology* **17**:2882-2892.

93 McGovern, S., C. D. Evans, P. Dennis, C. Walmsley, and M. A. McDonald. 2011. Identifying drivers of
94 species compositional change in a semi-natural upland grassland over a 40-year period.
95 *Journal of Vegetation Science* **22**:346-356.

96 Paulissen, M. P. C. P., P. J. M. van der Ven, A. J. Dees, and R. Bobbink. 2004. Differential effects of
97 nitrate and ammonium on three fen bryophyte species in relation to pollutant nitrogen
98 input. *New Phytologist* **164**:451-458.

99 Payne, R. J., N. B. Dise, C. J. Stevens, D. J. Gowing, and B. Partners. 2013. Impact of nitrogen
100 deposition at the species level. *Proceedings of the National Academy of Sciences of the*
101 *United States of America* **110**:984-987.

102 Provoost, S., M. L. M. Jones, and S. E. Edmondson. 2011. Changes in landscape and vegetation of
103 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 Remke, E., E. Brouwer, A. Kooijman, I. Blindow, and J. G. M. Roelofs. 2009. Low Atmospheric
107 Nitrogen Loads Lead to Grass Encroachment in Coastal Dunes, but Only on Acid Soils.
108 *Ecosystems* **12**:1173-1188.

109 Roem, W. J. and F. Berendse. 2000. Soil acidity and nutrient supply ratio as possible factors
110 determining changes in plant species diversity in grassland and heathland communities.
111 *Biological Conservation* **92**:151-161.

112 RoTAP. 2012. Review of Transboundary Air Pollution (RoTAP): Acidification, Eutrophication, Ground
113 Level Ozone and Heavy Metals in the UK Centre for Ecology & Hydrology, Edinburgh.

114 Seed, L., P. Wolseley, L. Gosling, L. Davies, and S. A. Power. 2013. Modelling relationships between
115 lichen bioindicators, air quality and climate on a national scale: Results from the UK OPAL air
116 survey. *Environmental Pollution* **182**:437-447.

117 Sheppard, L. J., I. D. Leith, T. Mizunuma, J. N. Cape, A. Crossley, S. Leeson, M. A. Sutton, N. van Dijk,
118 and D. Fowler. 2011. Dry deposition of ammonia gas drives species change faster than wet
119 deposition of ammonium ions: evidence from a long-term field manipulation. *Global Change*
120 *Biology* **17**:3589-3607.

121 Sheppard, L. J., I. D. Leith, T. Mizunuma, S. Leeson, S. Kivimaki, J. N. Cape, N. van Dijk, D. Leaver, M.
122 A. Sutton, D. Fowler, L. J. L. Van den Berg, A. Crossley, C. Field, and S. Smart. 2014. Inertia in
123 an ombrotrophic bog ecosystem in response to 9 years' realistic perturbation by wet
124 deposition of nitrogen, separated by form. *Global Change Biology* **20**:566-580.

125 Smart, S. M., M. R. Ashmore, M. Hornung, W. A. Scott, D. A. Fowler, U. Dragosits, D. C. Howard, M. A.
126 Sutton, and D. Famulari. 2004. Detecting the Signal of Atmospheric N Deposition in Recent
127 National-Scale Vegetation Change Across Britain. *Water, Air, & Soil Pollution: Focus* **4**:269-
128 278.

129 Smart, S. M., P. A. Henrys, B. V. Purse, J. M. Murphy, M. J. Bailey, and R. H. Marrs. 2012. Clarity or
130 confusion? - Problems in attributing large-scale ecological changes to anthropogenic drivers.
131 *Ecological Indicators* **20**:51-56.

132 Smart, S. M. and W. A. Scott. 2004. Bias in Ellenberg indicator values - problems with detection of
133 the effect of vegetation type. *Journal of Vegetation Science* **15**:843-846.

134 Southon, G. E., C. Field, S. J. M. Caporn, A. J. Britton, and S. A. Power. 2013. Nitrogen Deposition
135 Reduces Plant Diversity and Alters Ecosystem Functioning: Field-Scale Evidence from a
136 Nationwide Survey of UK Heathlands. *Plos One* **8**.

137 Stevens, C. J., N. B. Dise, D. J. G. Gowing, and J. O. Mountford. 2006. Loss of forb diversity in relation
138 to nitrogen deposition in the UK: regional trends and potential controls. *Global Change*
139 *Biology* **12**:1823-1833.

140 Stevens, C. J., N. B. Dise, J. O. Mountford, and D. J. Gowing. 2004. Impact of nitrogen deposition on
141 the species richness of grasslands. *Science* **303**:1876-1879.

142 Stevens, C. J., C. Dupre, E. Dorland, C. Gaudnik, D. J. G. Gowing, A. Bleeker, M. Diekmann, D. Alard, R.
143 Bobbink, D. Fowler, E. Corcket, J. O. Mountford, V. Vandvik, P. A. Aarrestad, S. Muller, and N.
144 B. Dise. 2010a. Nitrogen deposition threatens species richness of grasslands across Europe.
145 *Environmental Pollution* **158**:2940-2945.

146 Stevens, C. J., P. Manning, L. J. L. van den Berg, M. C. C. de Graaf, G. W. W. Wamelink, A. W. Boxman,
147 A. Bleeker, P. Vergeer, M. Arroniz-Crespo, J. Limpens, L. P. M. Lamers, R. Bobbink, and E.
148 Dorland. 2011. Ecosystem responses to reduced and oxidised nitrogen inputs in European
149 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* **166**: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 **154**: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, G.-R. 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.

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1 **Figure legends**

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⁻¹y⁻¹.

5 **Figure 2** Effect sizes (slopes) of NH_x:NO_y ratio in deposition on the number of species for UK
6 habitats. Only significant effects are shown. Actual ranges of NH_x:NO_y 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
10 vegetation type differs but slopes are shown for a range between 5 and 45 kgNha⁻¹y⁻¹.

11 **Figure 4** Effect sizes (slopes) of NH_x:NO_y ratio in deposition on the Ellenberg N number for
12 UK habitats. Only significant effects are shown. Actual ranges of NH_x:NO_y ratio covered by
13 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.
15 Only significant effects are shown. Actual ranges of N deposition covered by each vegetation
16 type differs but slopes are shown for a range between 5 and 45 kgNha⁻¹y⁻¹.

17 **Figure 6** Effect sizes (slopes) of NH_x:NO_y ratio in deposition on the Grass:Forb ratio for UK
18 habitats. Only significant effects are shown. Actual ranges of NH_x:NO_y ratio covered by each
19 vegetation type differs but slopes are shown for a range between 0.5 and 3.5.

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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_x or NO_y with respectively NO_y and NH_x 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.

			<i>Sp. rich</i>	<i>Ellenberg N</i>	<i>Grass:Forb</i>
Dry upland heaths	A	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*
		NH _x :NO _y ratio	0.783	0.475***	0.528
	B	NO _y deposition	-0.037	-0.007	-0.001
		NH _x deposition	-0.019	0.009	0.012
Dry lowland heaths	A	Temperature	0.474	0.103**	0.035
		Precipitation	0.010	-0.002**	0.005*
		S deposition ¹	-0.320	-0.480	1.290
		Total N deposition	-0.266***	-0.022***	0.035*
		NH _x :NO _y ratio	1.148	0.634***	0.342
	B	NO _y deposition	-0.099	-0.021*	0.001
		NH _x deposition	-0.030	0.006	0.011
Bogs and Mires	A	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***
		NH _x :NO _y ratio	0.797	0.374***	0.312
	B	NO _y deposition	-0.044	-0.006	0.000
		NH _x deposition	0.004	0.007*	0.009
Base rich Mires	A	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
		NH _x :NO _y ratio	-0.364	-0.046	0.081
	B	NO _y deposition	0.004	0.004	0.003
		NH _x deposition	-0.014	0.008	-0.001

6 Number of plots: Dry upland heaths (267), dry lowland heaths (182), Bogs and Mires (1136), Base rich Mires (274). ¹ S deposition was
7 inverse transformed

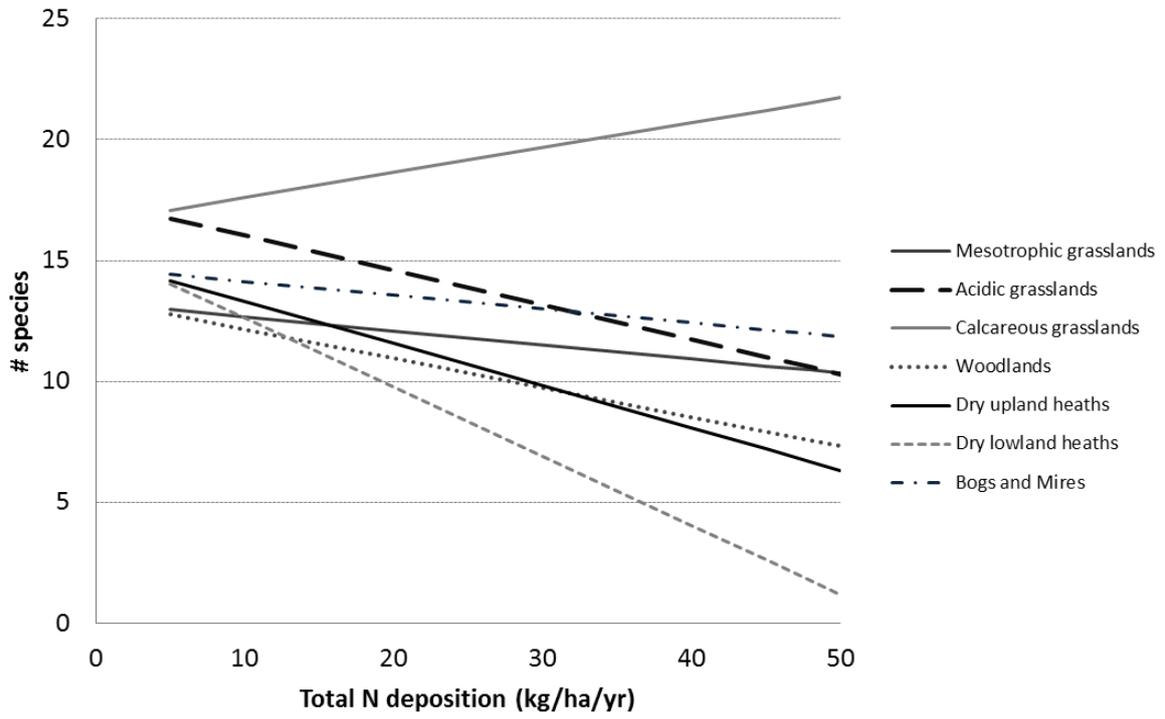
8 **Table 2** Coefficients and their significance for the habitats acidic grasslands, mesotrophic grasslands,
 9 calcareous grassland and woodlands. Results of a multiple regression (A) and regression analysis of
 10 the residuals of models containing either NH_x or NO_y with respectively NO_y and NH_x to separate N
 11 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.

			<i>Sp. rich</i>	<i>Ellenberg N</i>	<i>Grass:Forb</i>
Acidic grasslands	A	Temperature	0.097	0.086***	-0.098**
		Precipitation ¹	-0.003***	0.016***	-0.040
		S deposition	-0.113**	0.005	-0.013
		Total N deposition	-0.145***	-0.015***	0.030***
		NH _x :NO _y ratio	-1.783***	0.368***	0.455**
	B	NO _y deposition	-0.011	-0.018***	0.002
		NH _x deposition	-0.061*	0.006	-0.001
Mesotrophic grasslands	A	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
		NH _x :NO _y ratio	-0.947**	0.172***	0.100
	B	NO _y deposition	0.023	-0.010	-0.001
		NH _x deposition	-0.096***	0.018***	0.003
Calcareous grasslands	A	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***
		NH _x :NO _y ratio	-2.849***	0.618***	0.258*
	B	NO _y deposition	0.259***	-0.025**	-0.026
		NH _x deposition	-0.109**	0.046***	-0.002
Woodlands	A	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
		NH _x :NO _y ratio	1.178*	0.271**	-0.116
	B	NO _y deposition	-0.229***	-0.022	0.017
		NH _x deposition	0.019	0.026***	-0.015

13 *Number of plots: Acidic grasslands (1090), Mesotrophic grasslands (1195), Calcareous grasslands (830), Woodlands (514).* ¹ *Precipitation*
 14 *was inverse transformed*

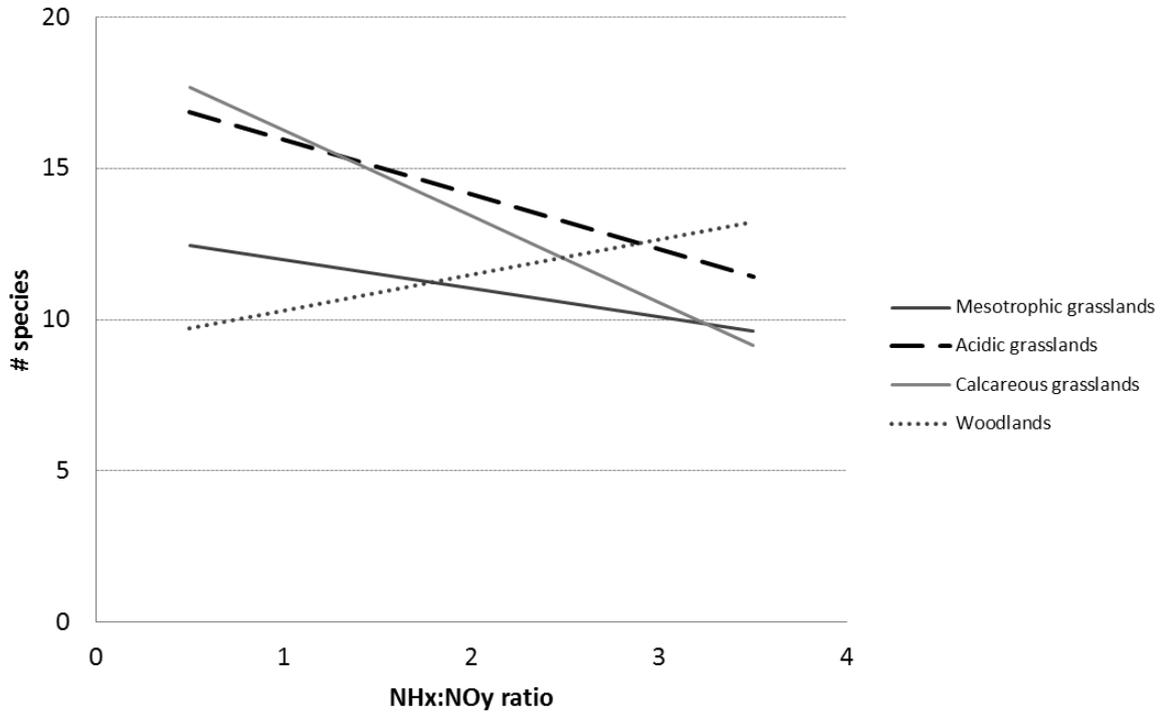
15

1 **Figure 1**



2

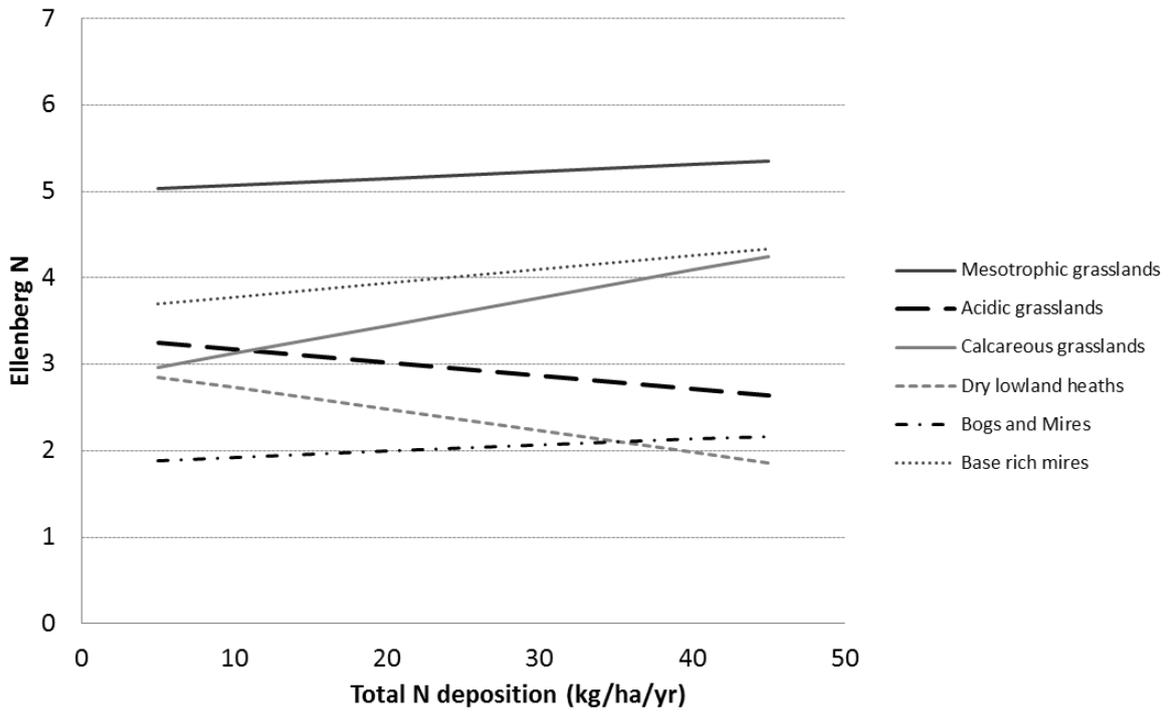
3 **Figure 2**



4

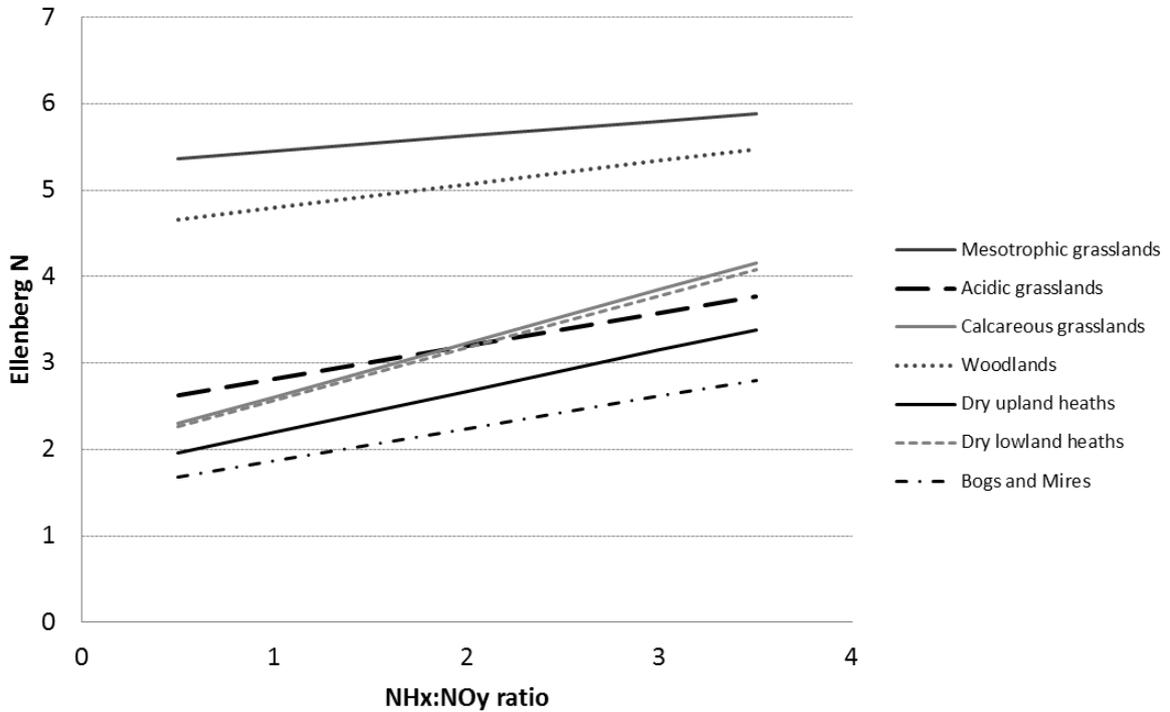
5

6 **Figure 3**



7

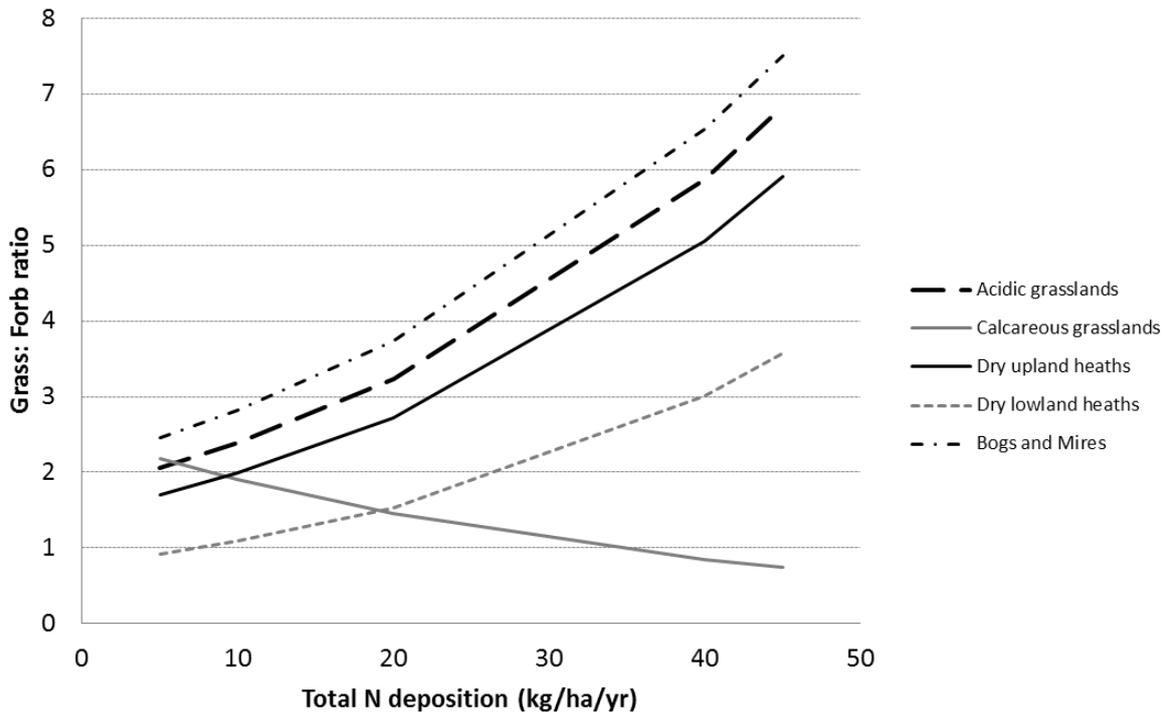
8 **Figure 4**



9

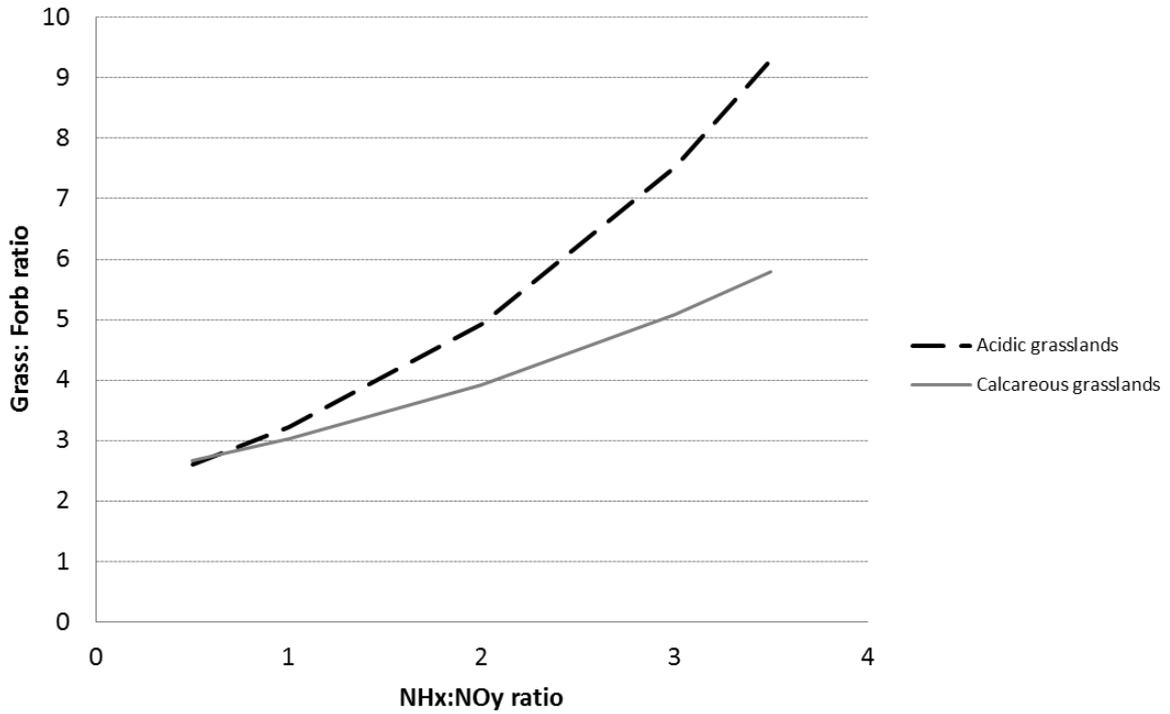
10

11 **Figure 5**



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13 **Figure 6**



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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

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