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Plant invasion is associated with higher plant-soil nutrient concentrations in nutrient poor-environments

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Abstract

Plant invasion is an emerging driver of global change worldwide. We aimed to disentangle its impacts on plant-soil nutrient concentrations. We conducted a meta-analysis of 215 peer-reviewed articles and 1233 observations. Invasive plant species had globally higher N and P concentrations in photosynthetic tissues but not in foliar litter, in comparison to their native This article has been accepted for publication and undergone full peer review but has not been through

the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as doi: 10.1111/gcb.13384 This article is protected by copyright. All rights reserved. competitors. Invasive plants were also associated with higher soil C and N stocks and N, P and K availabilities. The differences in N and P concentrations in photosynthetic tissues and in soil total C and N, soil N, P and K availabilities between invasive and native species decreased when the environment was richer in nutrient resources. The results thus suggested higher nutrient resorption efficiencies in invasive than in native species in nutrient-poor environments. There were differences in soil total N concentrations but not in total P concentrations, indicating that the differences associated to invasive plants were related with biological processes, not with geochemical processes. The results suggest that invasiveness is not only a driver of changes in ecosystem species composition but that it is also associated with significant changes in plant-soil elemental composition and stoichiometry.

Keywords: C:N, soil fertility, N:P, nitrogen, phosphorus, potassium

Introduction

The structure, diversity and production capacity of terrestrial ecosystems is strongly linked to the concentrations and stoichiometric ratios in the different ecosystemic compartments and the soil availability of nitrogen, phosphorus and potassium (Sterner & Elser, 2002; Reich & Oleksyn, 2004; Elser *et al.*, 2007; Vitousek *et al.*, 2010; Sardans *et al.*, 2011; Peñuelas *et al.*, 2013; Sardans & Peñuelas, 2013). Most drivers of global change, such as increasing atmospheric CO₂ concentrations, N eutrophication, drought, warming or land-use changes change those elemental compositions and stoichiometries of ecosystemic compartments and those relationships with ecological processes and species composition (Seabloom *et al.*, 2006; Elser *et al.*, 2010; Sardans & Peñuelas, 2012; Sardans *et al.*, 2012; Peñuelas *et al.*, 2013; Yuan & Chen, 2015). The growing success of invasive plants in many regions; 20% or more of plant

species are exotics in many continental areas and 50% or more in islands (Seabloom *et al.*, 2006), e.g. plant invaders are affecting 405,000 Km² in United States (Seabloom *et al.*, 2015) is an emerging driver of Global Changes; however, it has not received the same level of attention at this regard of the impacts on plant-soil nutrient concentrations (Hulme *et al.*, 2009, 2015).

Previous studies have observed that several mechanisms involved in the uptake and nutrient use efficiency by plants underlie the success of invasive plants (Daehler, 2003; González et al., 2010). The mechanisms seem to differ between nutrient-poor and nutrient-rich soils. In nutrient-poor soils most studies suggest that the success of invasive plants depends on conservative strategies, such as a higher nutrient-use efficiency (Ostertag & Verville, 2002; Funk & Vitousek, 2007; González et al., 2010; Matzek, 2011), especially on short time scales (Funk & Vitousek, 2007), long nutrient residence times (Laungani & Knops, 2009), high resistance to low levels of nutrients (Kueffer, 2009; Schumacher et al., 2000) and high plasticity of stoichiometric ratios (González et al., 2010). In fact, all these traits are consistent with those expected in stress tolerant species (Grime 1977), in this case by a stress due to nutrient limitation. The establishment of new symbiosis (Hiltbrunner et al., 2014) or the more effective use of existing symbiosis (Pringle et al., 2009) are other strategies frequently linked to plant invasiveness success, all them increasing the availability of limiting soil resources. In contrast, in nutrient-rich soils, there is an advantage of species with high rates of photosynthesis and growth (Schumacher et al., 2000; González et al., 2010), high reproductive outputs (González et al., 2010), large body size (Van Kleunen et al., 2010), low C:nutrient ratios in tissues (Schumacher et al., 2000; Peñuelas et al., 2009; González et al., 2010), low costs of foliar construction (Nagel & Griffin, 2001; González et al., 2010), large investments of N in photosynthetic production (Ehrenfeld, 2003; Shen et al., 2011), high capacities of nutrient uptake (Zabinski et al., 2002; Leffler et al., 2011; Peng et al., 2011) and high levels of plasticity in the acquisition of resources as a function of pulses in nutrient availability (Leffler et al., 2011).

Nutrient uptake and all foliar traits enabling rapid rates of growth (Zabinski *et al.*, 2002; Leihsman *et al.*, 2007) will thus help invading species to succeed when resources are not limited (Leihsman *et al.*, 2007; Peng *et al.*, 2011). Some authors have claimed that, independently of growth conditions, invaders are more likely to have higher foliar areas, lower tissue construction costs and greater phenotypical plasticity that increase the availability of soil resources (Daehler, 2003).

Invasive-plant success has also been linked to differences in soil elemental composition. In a recent review, Pysek *et al.* (2012) reported that 192 of 436 case studies on the effects of invasive plants on soil nutrient concentrations found higher concentrations, 72 found lower concentrations and 158 found no significant differences. Sardans & Peñuelas (2012), by analyzing 65 case studies, showed that most processes of invasion had higher availability of soil nutrients. In addition to these previous qualitative studies, Vila *et al.* (2011) conducted a metaanalysis on the relationships of plant invasive success with soil condition showing that invasive success is related with higher soil C, N and P stocks.

There is, however, no general consensus on whether or not successful plant invaders have different elemental compositions than the native species, or, if present, whether differences are dependent or not on habitat nutrient richness. A quantitative study comparing plant, litter and soil nutrient concentrations, i.e. the whole plant-soil system, between invasive and their native competitors at the global scale is missing. Moreover, there are no studies analyzing the differences for other important elements, such as K. In addition to the possible influence of soil nutrient-richness, the possible influence of climate conditions on these relationships warrants investigation since climatic shifts affect invasive plant functional processes and in general invasion patterns (Lu *et al.*, 2013; Zenni & Hoban, 2015), and thus could affect the differences in plant, litter and soil nutrient concentrations between invasive and

native plants at the global scale. In regions where climate evolves towards characteristics more favorable to plant production (higher MAP and/or MAT) and where invasive success is expected to be related to higher rates of nutrient-uptake and in general to C and/or R ecological strategies (Grime, 1977), we should expect more investment of nutrients in plant growth and faster nutrient cycling rates in plant-soil system. Contrarily, in regions evolving towards more extreme and stressed climatic conditions, we should expect invassive success to be related to more conservative traits, less growth, traits typical of stress-tolerator biological strategy (T strategy, Grime 1977) that are less linked with higher-uptake capacity, but to a higher resorption and retention of nutrients in the system and consequently with higher nutrient concentrations in plant-soil system.

We have conducted a global meta-analysis of both the past and the most recent literature data on the nutrient concentrations in photosynthetic tissues, foliar litter and soil with the aims to determine whether or not invasive-plant success (i) is associated with different elemental compositions of photosynthetic tissues and foliar litter between successful invasive plants and their native competitors, (ii) is associated with changes in soil elemental composition and nutrient availability and stoichiometry, and (iii) how these associations, if exist, depend on soil nutrient concentrations and availabilities, and climatic condition.

Materials and methods

Data collection

We searched the ISI Web of Science using combinations of the following keywords: alien, availability, available, carbon, concentration, C:K, C:N, C:P, foliar, invasion, invasive, leaf, needle, nitrogen, N:K, N:P, phosphorus, plant, potassium, P:K, ratio, soil, solution, stoichiometric, stoichiometry, success. We only selected studies providing the same equivalent information for invasive successful species and their native competitors. Moreover, we only analyzed plant, litter and soil variables with a minimum of 45

different reports that included the information for invasive and the respective native species. These variables finally included N and P concentration and C:N concentration ratio in photosynthetic tissues, foliar-litter N concentration and C:N concentration ratio, and soil total C, N and P concentrations, total soil C:N concentration ratio, soil P-Olsen and soil extractable K⁺, NO₃⁻ and NH₄⁺ concentrations. In the few studies with different temporal data we used the average mean values. Finally, only field non manipulative studies have been considered. Applying these criteria we obtained 215 reports with 1233 observations across the world (Figure S1).

Climatic data

We extracted climatic data for each study site from the WorldClim database (Hijmans *et al.*, 2005). This database provides global maps of interpolated variables of climatological variables extrapolated from extensive climatic time series (from 1950 to 2000), with a spatial resolution of 30 arc seconds (~1 km at the equator). We used MAT and MAP as climatic predictor variables.

Data analyses

We examined the effects of invasive-plant success on the differences of photosynthetic tissues elemental compositions and stoichiometries and soil nutrient status between successful invasive plants and their native competitors by calculating the ln response ratios from each study as described by Hedges *et al.* (1999). The natural ln response -ratio (ln*RR*) was calculated as ln (X/X_n) = ln X_i – ln X_n , where X_i and X_n are the values of each observation in the invaded soil or invasive plant and in the corresponding native situation, respectively. The sampling variance for each ln*RR* was calculated as ln[($1/n_i$) × (S_i/X_i)² + ($1/n_n$) × (S_n/X_n)²] using the R package metafor 1.9–2, where n_i , n_n , S_n , S_n , X_i and X_n are the invasive and native sample sizes, standard deviations, and mean response values, respectively. The natural ln response ratios were determined by specifying studies as random factors using the *rma* model in metafor. The effects on soil elemental variables and the difference between the elemental compositions of invasive and native plants were considered significant if the 95% confidence interval (CI) of ln*RR* did not overlap zero. All these statistical analyses were performed in R 3.1.2 (R Core Team, 2015). Despite for most studied variables there was a low proportion of studies containing N₂-fixing species, we performed these analyses

twice, once with the entire data another one with and after the removal of the studies that contained N_2 fixing plant species for detecting the possible importance of N_2 -fixing capacity in the ln response ratio effect of the plant and soil variables studied. We analyzed variables with more than 45 observations available at the global scale. The number of reports and observations used by statistical analyses of each studied soil, plant and litter variable are shown in Figures 1 and 2, and described in Tables S1-S3.

We also examined whether the differences in the In response ratio of plants and soils depend on environmental circumstances such as climate or soil total nutrient concentration and soil available nutrient concentration. For these analyses, we related the In response ratio effect mentioned above (InRR) with climatic variables at each study site. We used MAP and MAT data from the WorldClim database (Hijmans et al., 2005). We also tested whether InRR is dependent on native plant and soil total nutrient concentrations and soil nutrient availability. We conducted a regression of the In response ratio of the soil N concentration (In invaded soil N value - In native soil N value) relative to the concentration in the natural (native site) soil (In native soil N value). In the case of foliar plant tissues, nutrient concentration has generally been well correlated with soil nutrient availability across natural gradients or fertilization experiments (Porder et al., 2005; Alvarez-Clare & Mack, 2015). We thus used the native foliar concentrations as a proxy of site soil availability to relate the possible differences in the In response ratio effect in foliar and foliar-litter variables (In invasive plant N value - In native plant N value), with the corresponding variable availability in soil (In native foliar N value). We used regression type II for these analyses, because both dependent and independent variables were interchangeable and random, so the error of the independent variable could not be neglected. We ran a standardized major axis method (SMA) using the SMATR package (Warton et al., 2006) (http://www.bio.mq.edu.au/ecology/SMATR).

Finally, in the cases of total soil N concentration, soil P-Olsen and foliar N and P concentrations, for which we have the larger number of observations, we divided the observations of each one of these variables according with their values in native soils or plants in three groups with similar number of observations. Thus, the groups corresponded to low, intermediate and high values in native conditions as

a proxy of site nutrient richness. Thereafter we conducted an one-way ANOVA with Bonferroni post-hoc test to detect possible differences in the In response ratio among the three groups.

Results

Differences in photosynthetic tissues and foliar litter

A meta-analysis of the entire data set indicated that invasive plant species had higher N (z = 8.93, P < 0.0001) and P (z = 3.44, P < 0.001) concentrations (41% and 32%, respectively) and lower (26%) C:N ratios (z = - 5.02, P < 0.0001) in their photosynthetic tissues than the native competitors (Fig. 1a). An analysis of the same data set but without excluding N₂-fixing species also indicated higher N (z = 6.57, P < 0.0001) and P (z = 2.67, P < 0.01) concentrations (29% and 32%, respectively) and lower (22%) C:N ratios (z = - 4.84, P < 0.0001) in the photosynthetic tissues of the invasive species (Fig. 1b). The N concentration and the C:N ratio in foliar litter were, however, not significantly different either for the entire data set (Fig. 1c) or when the data for the N₂-fixing plant species were excluded (Fig. 1d). Not significant differences were either found for litter P concentrations (only 13 observations, data not shown).

Differences in soil conditions

The soil concentrations of extractable K (z = 2.53, P < 0.05), soluble nitrate (z = 7.40, P < 0.0001), P-Olsen (z = 2.83, P < 0.01) and total N (z = 4.34, P < 0.0001) and C concentrations (z = 3.62, P < 0.001) were higher (13%, 117%, 21%, 19% and 12%, respectively) in soils of invasive plants than in soils of their corresponding native competitor species. The concentration of soluble ammonium was also marginally (z = 1.81, P = 0.07) higher (11%) in the soils of the invasive than the native species. The ln response ratio effects on the soil C:N ratio and total P concentration were not statistically significant.

An analysis of the same data set but without the data for the N₂-fixing species produced similar results (Fig. 2b). The soluble nitrate (z = 6.37, P < 0.0001), P-Olsen (z = 2.83, P < 0.001), total N (z = 2.32, P < 0.05) and C (z = 3.13, P < 0.001) concentrations, were higher (118%, 27%, 10% and 7%, respectively) in the soils of the invasive plants than in the soils of the native competitors. The concentration of extractable K was marginally (z = 1.80, P = 0.072) higher (11%) in the soils of the invasive species.

Ln response ratios along gradients of nutrient availability and climate

The In response ratio of total N and P concentrations in photosynthetic tissues of invasive plants were negatively correlated to the corresponding values for the photosynthetic tissues of the native plant competitors (Figs. 3a and 3b). The In response ratio of foliar N concentration was positively different from zero in sites with low and intermediate values, whereas for foliar P concentrations the In response ratio was only positively different from zero in sites with low values (Figs. S2a and S2b). No significant relationships were observed between foliar litter N and P In response ratio and the corresponding values for the foliar litter of the native plant competitors (Figs. 3c and 3d).

The In response ratio for soil total N, P-Olsen, soluble nitrate and extractable K concentrations in invaded soils were negatively correlated with the corresponding values in the soils of the native plant competitors (Figs. 4a–d). For soil nitrate concentration, total N concentration and soil P-Olsen, the In response ratio was positively different than zero in sites with low and intermediate values, whereas for soil available K⁺ the In response ratio was positively different than zero only in sites with low values (Fig. S3).

Interestingly, few relationships between climatic gradients and ln response-ratio effects were detected. MAT was positively but weakly correlated with the ln response ratios for soil total N concentration (R = 0.27, P < 0.001) and with N concentration in photosynthetic tissues (R = 0.16, P < 0.05). MAP was positively and also weakly correlated with the ln response ratio for soil soluble nitrate concentration (R = 0.25, P < 0.01) (Fig. S4).

Discussion

Our study showed higher N and P concentrations in the photosynthetic tissues of invasive species in nutrient-poor environments. These higher concentrations were found in photosynthetic tissues but not in foliar litter, suggesting a higher N and P resorption capacity in resource-poor than in resource-rich environments. These results are consistent with previous studies observing that the competitive advantage over native plant species competitors and the success of invasive plants in resource-poor environments has frequently been correlated with a more conservative use of nutrients, higher residence time due to higher nutrient-resorption capacities (Ostertag & Verville, 2002), and higher photosynthetic nitrogen use efficiency (Ens *et al.*, 2015).

The soils under the invasive plants had higher soil P-Olsen, soluble nitrate and potassium concentrations and therefore higher availability of the three most important soil macronutrients for plant growth. The higher soil NO₃⁻ concentrations in soils under invasive species than under their native competitors is consistent with previous studies observing a positive relationship between soil NO₃⁻ concentration and the intensity of plant species invasive success (Gilliam, 2006). The studies compiled in this meta-analysis did not allow a clear determination of whether these higher concentrations were the cause or the effect of the success of invasive plant species. The studies that have experimentally tested whether soil

differences were the cause or the consequence of plant invasion, however, have reported that soil differences were mainly due to the effect of the success of the invasive species (Li et al., 2006; Dassonville et al., 2008; Elgersma et al., 2011; Lee et al., 2012; Kuedding et al., 2014; Stark & Norton, 2014). A few number of reports that have studied the changes in soil conditions during 4 (Belnap et al., 2005) and 7 (Hawkes et al., 2005) years have observed that the invasive species changed soil conditions over time. Several studies have also observed a direct impact of invasive-plant establishment on soil function such as increases in soil enzymatic activities associated with increases in some soil elemental concentrations (Hawkes et al., 2005; Alison et al., 2006; Caldwell, 2006; Aragon et al., 2014; Kuebbing et al., 2014), mineralization (Haubensak & Parker, 2004; Fickbohm & Zhu, 2006; Li et al., 2006) and respiration (Souza-Alonso et al., 2015). Other studies, although fewer than the above, did not observe these differences in soil enzymatic activity and mineralization (Zabinsky et al., 2002, Meisner et al., 2011) or found different results depending on species and site (Koutika et al., 2007) or on the enzymatic activities (Chacón et al., 2009). Kulmatiski et al. (2006) in 660 experimental plots in abandoned croplands (from 50 to 7 years ago) with different management histories observed that invasive success explained the soil C, N and P concentrations more significantly than the previous agricultural histories, suggesting that the invasive plants facilitated their own growth by maintaining beneficial fungal communities and fast nutrient-cycling rates.

Our results showed a general globally higher soil total N concentration under invasiveplants than under their native competitors but we did not observe a higher total P concentration. Cycling and concentrations of soil N mainly depends on biological processes, whereas mineral rocks are the sources of soil P, and its soil total concentration is primarily driven by physicogeochemical processes (Gómez-Aparicio & Canhan, 2008; Vitousek *et al.*, 2010; Peñuelas *et al.*, 2013). Both N and P are important soil components that could be involved in facilitating plant invasion, but only soil total N concentration can thus be associated mostly with

the biological process of plant invasion. This fact is consistent with the hypothesis that the differences between the soils under invasive and native plants are most likely due to the effects of species invasion itself. Rather surprisingly, the effects of plant-invasions on soil and plant N concentrations, C:N ratios and most other significant stoichiometry parameters were not different when including N₂-fixing plants than when excluding them from the global analysis. Changes in soil physical conditions or in microbial communities, including soil N-fixing microbes, could be involved, warranting further research.

The differences in soil total C and N, and in N, P and K availabilities and in N and P concentrations in photosynthetic tissues between invasive and native species decreased with increasing values of the corresponding variables in natural-native conditions to the point that the differences disappeared in resource-rich environments. These lower differences in resource-rich environments could be due to the higher nutrient up-take in invasive species being counteracted by its higher growth capacity, and the corresponding dilution effect. In nutrient-rich sites, moreover, native species are also highly competitive, having traits that enable native plants to be very effective in taking up resources. On the other hand, the frequently observed higher mineralization capacity and enzyme activity under invasive than under native species in nutrient-rich soils (Allison *et al.*, 2006; Gómez-Aparicio & Canham, 2008; Aragón *et al.*, 2014) would increase the rates of nutrient released from organic matter, but this would be also counteracted by the higher plant nutrient uptake so that soil nutrient concentrations would remain similar than under native species.

Climate variables had few correlations with the studied In response ratio of the studied variables. MAT had weak but positive correlation with In response ratio of total soil N concentration and N concentration in plant tissues and MAP had also a positive relationship with soil nitrate In response ratio. These results suggest thus that climatic conditions are less

influential on the In response ratio of the studied soil total and available nutrient concentrations than the environmental nutrient richness.

Summarizing, this is the first study that has analyzed globally the association between plant invasion and nutrient concentration and stoichiometry of photosynthetic tissues, leaf litter and soils. Invasive plant species had globally higher N and P concentrations in photosynthetic tissues but not in foliar litter, in comparison to their native competitors. Invasive plants were also associated with higher soil C and N stocks and N, P and K availabilities. The differences in N and P concentrations in photosynthetic tissues and in soil total C and N, soil N, P and K availabilities between invasive and native species decreased when the environment was richer in nutrient resources. These global trends may be explained by (i) larger differences in resorption and nutrient-use efficiency between invasive and native species in nutrient-poor environments, and (ii) a higher competitive capacity associated with larger nutrient uptake and plant growth capacity with a dilution effect in invasive than in native species when environments become richer in resources. Moreover, some other mechanisms such as enhancement of soil enzymatic activity and mineralization, and more effective symbiotic relationships can be also involved in these global trends. Clearly determining whether invasive-plant success is the cause or the consequence of soil elemental composition and nutrient availability is currently not possible, but research up to now suggests that these plant and soil nutritional changes are more the consequence than the cause of plant invasion. Plant invasiveness should thus not be neglected as a driver of global change in plant-soil elemental and stoichiometric composition and soil fertility.

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References

- Allison SD, Nielsen C, Hughes RF (2006) Elevated enzyme activities in soils under the invasive nitrogenfixing tree *Falcataria moluccana*. Soil Biology and Biochemistry **38**, 1537–1544.
- Alvarez-Clare S, Mack MC (2015) Do Foliar, Litter, and Root Nitrogen and Phosphorus Concentrations Reflect Nutrient Limitation in a Lowland Tropical Wet Forest? *PLoS One* **10**, e0123796.
- Aragón R, Sardans J, Peñuelas J (2014) Soil enzymes associated with carbon and nitrogen cycling in invaded and native secondary forests of northwestern Argentina. *Plant Soil* **384**, 169–183.
- Belnap J, Phillips SL, Sherrod SK, Moldenke A (2005) Soil biota can change after exotic plant invasion: does this affect ecosystem processes? *Ecology* **86**, 3007–3017.
- Caldwell BA (2006) Effects of invasive scotch broom on soil properties in a Pacific coastal prairie soil. Applied Soil Ecology **32**, 149–152.
- Chacón N, Herrera I, Flores S, González JA, Nassar JM (2009) Chemical, physical, and biochemical soil properties and plant roots as affected by native and exotic plants in Neotropical arid zones. *Biology and Fertility Soils* **45**, 321–328.
- Daehler CC (2003) Performance Comparisons of Co-Occurring Native and Alien Invasive Plants: Implications for Conservation and Restoration. *Annual Review of Ecology Evolution and Systematics* **34**, 183–211.
- Dassonville N, Vanderhoeven S, Vanparys V, Hayez M, Gruber W, Meerts P (2008) Impacts of alien invasive plants on soil nutrients are correlated with initial site conditions in NW Europe. *Oecologia* 157, 131–140.
- Ehrenfeld JG (2003) Effects of Exotic Plant Invasions on Soil Nutrient Cycling Processes. *Ecosystems* **6**, 503–523.
- Elgersma KJ, Ehrenfeld JG, Yu S, Vor T (2011) Legacy effects overwhelm the short-term effects of exotic plant invasion and restoration on soil microbial community structure, enzyme activities, and nitrogen cycling. *Oecologia* **167**, 733–745.
- Elser JJ, Fagan WF, Kerkhoff A, Swenson NG, Enquist B (2010) Biological stoichiometry of plant production: metabolism, scaling and ecological response to global change. *New Phytologist* **186**, 593–608.
- Elser JJ, Bracken MES, Cleland EE, *et al.* (2007) Global analysis of nitrogen and phosphorus limitation of primary producers in freshwater, marine and terrestrial ecosystems. *Ecology Letters* **10**, 1135–1142.
- Ens E, Hutley LB, Rossiter-Rachor NA, Douglas MM, Setterfield SA (2015) Resource-use efficiency explains grassy weed invasion in a low-resource savanna in north Australia. *Frontieres in Plant Science* **6**, 50.
- Fickbohm SS, Zhu WX (2006) Exotic purple loosestrife invasion of native cattail freshwater wetlands: Effects on organic matter distribution and soil nitrogen cycling. *Applied Soil Ecology* **32**, 123–131.
- Funk JL, Vitousek PM (2007) Resource-use efficiency and plant invasion in low-resource systems. *Nature* **446**, 1079–1081.
- Gilliam FS (2006) Response of the herbaceous layer of forest ecosystems to excess nitrogen deposition. *Journal of Ecology* **94**, 1176-1191.
- Gómez-Aparicio L, Canham CD (2008) Neighborhood Models of the Effects of Invasive Tree Species on Ecosystem Processes. *Ecological Monographs* **78**, 69–86.
- González AL, Kominoski JS, Danger M, Ishida S, Iwai N, Rubach A (2010) Can ecological stoichiometry help explain patterns of biological invasions? *Oikos* **119**, 779–790.

Haubensak KA, Parker IM (2004) Soil changes accompanying invasion of the exotic shrub *Cytisus scoparius* in glacial outwash prairies of western Washington [USA]. *Plant Ecology* **175**, 71–79.

Grime JP (1977) Evidence for existence of three primary strategies in plants and its relevance to ecological and evolutionary theory. *American Naturalist* **982** 1169-1194.

- Hawkes CV, Wren IF, Herman DJ, Firestone MK (2005) Plant invasion alters nitrogen cycling by modifying the soil nitrifying community. *Ecology Letters* **8**, 976–985.
- Hedges LV, Gurevitch J, Curtis PS (1999) The meta-analysis of response ratios in experimental ecology. *Ecology* **80**, 1150–1156.
- Hijmans RJ, Cameron SE, Parra JL, Jones PG, Jarvis A (2005) Very high resolution interpolated climate surfaces for global land areas. *International Journal of Climatology* **25**, 1965–1978.
- Hiltbrunner E, Aerts R, Bühlmann T, *et al.*, (2014) Ecological consequences of the expansion of N₂-fixing plants in cold biomes. *Oecologia* **176**, 11–24.
- Hulme PE (2015) Invasion pathways at a crossroad: policy and research challenges for managing alien species introductions. *Journal of Applied Ecology* **52**, *1418-1424*.
- Hulme PE (2009) Trade, transport and trouble: Managing invasive species pathways in an era of globalization. *Journal of Applied Ecology* **46**, 10–18.
- Koutika LS, Vanderhoeven S, Chapuis-Lardy L, Dassonville N, Meerts P (2007) Assessment of changes in soil organic matter after invasion by exotic plant species. *Biology and Fertility of Soils* **44**, 331– 341.
- Kuebbing SE, Classen AT, Simberloff D (2014) Two co-occurring invasive woody shrubs alter soil properties and promote subdominant invasive species. *Journal of Applied Ecology* **51**, 124–133.
- Kueffer C (2009) Reduced risk for positive soil-feedback on seedling regeneration by invasive trees on a very nutrient-poor soil in Seychelles. *Biological Invasions* **12**, 97–102.
- Kulmatiski A, Beard KH, Stark JM (2006) Soil history as a primary control on plant invasion in abandoned agricultural fields. *Journal Applied Ecology* **43**, 868–876.
- Laungani, Knops JMH (2009) Species-driven changes in nitrogen cycling can provide a mechanism for plant invasions. *Proceedings of Nationall Academy of Sciences USA* **106**, 12400–12405.
- Lee MR, Flory SL, Phillips RP (2012) Positive feedbacks to growth of an invasive grass through alteration of nitrogen cycling. *Oecologia* **170**, 457–465.
- Leffler AJ, Monaco TA, James JJ (2011) Nitrogen acquisition by annual and perennial grass seedlings: Testing the roles of performance and plasticity to explain plant invasion. *Plant Ecology* **212**, 1601– 1611.
- Leishman MR, Haslehurst T, Ares A, Baruch Z (2007) Leaf trait relationships of native and invasive plants: Community- and global-scale comparisons. *New Phytologist* **176**, 635–643.
- Li WH, Zhang CB, Jiang HB, Xin GR, Yang ZY (2006) Changes in soil microbial community associated with invasion of the exotic weed, *Mikania micrantha* H.B.K. *Plant and Soil* **281**, 309–324.
- Lu X, Siemann E, Shao X, Wei H, Ding J (2013) Climate warming affects biological invasions by shifting interactions of plants and herbivores. *Global Change Biology* **19**, 2339–2347.
- Matzek V (2011) Superior performance and nutrient-use efficiency of invasive plants over non-invasive congeners in a resource-limited environment. *Biological Invasions* **13**, 3005–3014.
- Meisner A, de Boer W, Verhoeven KJF, Boschker HTS, van der Putten WH (2011) Comparison of nutrient acquisition in exotic plant species and congeneric natives. *Journal of Ecology* **99**, 1308–1315.
- Nagel JM, Griffin KL (2001) Construction cost and invasive potential: Comparing Lythrum salicaria (Lythraceae) with co-occurring native species along pond banks. American Journal of Botany 88, 2252–2258.
- Ostertag R, Verville JH (2002) Fertilization with nitrogen and phosphorus increases abundance of nonnative species in Hawaiian montane forests. *Plant Ecology* **162**, 77–90.
- Peng RH, Fang CM, Li B, Chen JK (2011) *Spartina alterniflora* invasion increases soil inorganic nitrogen pools through interactions with tidal subsidies in the Yangtze Estuary, China. *Oecologia* **165**, 797–807.
- Peñuelas J, Sardans J, Llusià J, *et al.*, (2009) Faster returns on 'leaf economics' and different biogeochemical niche in invasive compared with native plant species. *Global Change Biology* **16**, 2171–2185.
- Peñuelas J, Poulter B, Sardans J, *et al.*, (2013) Human-induced nitrogen–phosphorus imbalances alter natural and managed ecosystems across the globe. *Nature Communications* **4**, 2934.
- Pringle A, Bever JD, Gardes M, Parrent JL, Rillig MC, kilronomos JN (2009) Mycorrhizal Symbioses and

Plant Invasions. Annual Reviews in Ecology Evolution and Systematics 40, 699–715.

- Porder S, Asner GP, Vitousek PM (2005) Ground-based and remotely sensed nutrient availability across a tropical landscape. *Proceedings National Academy Sciences USA* **102**, 10909–10912.
- Pyšek P, Jarosik V, Hulme PE, Pergl J, Hejda M, Schaffner U, Vilà M (2012) A global assessment of invasive plant impacts on resident species, communities and ecosystems: the interaction of impact measures, invading species' traits and environment. *Global Change Biology* **18**, 1725–1737.
- Reich PB, Oleksyn J (2004) Global patterns of plant leaf N and P in relation to temperature and latitude. *Proceedings National Academy Sciences USA* **101**, 11001–11006.
- R Core Team (2015). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. URL https://www.R-project.org/.
- Sardans J, Rivas-Ubach A, Peñuelas J (2011) The elemental stoichiometry of aquatic and terrestrial ecosystems and its relationships with organismic lifestyle and ecosystem structure and function: a review and perspectives. *Biogeochemistry* **111**, 1–39.
- Sardans J, Penuelas J (2012) The Role of Plants in the Effects of Global Change on Nutrient Availability and Stoichiometry in the Plant-Soil System. *Plant Physiology* **160**, 1741–1761.
- Sardans J, Rivas-Ubach A, Peñuelas J (2012) The C:N:P stoichiometry of organisms and ecosystems in a changing world: A review and perspectives. *Perspectives Plant Ecology Evolution and Systematics* **14**, 33–47.
- Sardans J, Peñuelas J (2015) Potassium: a neglected nutrient in global change. *Global Ecology and Biogeography* **24**, 261–275.
- Seabloom EW, Williams JW, Slayback D, Stoms DM, Viers JH, Dobson AP (2006) Human impacts, plant invasion, and imperiled plant species in California. *Ecological Applications* **16**, 1338–1350.
- Seabloom, EW, Borer ET, Buckley YM, et al., (2015) Plant species' origin predicts dominance and response to nutrient enrichment and herbivores in global grasslands. *Nature Communications* **6**, 7710.
- Shen XY, Peng SL, Chen BM, Pang JX, Chen LY, Xu HM, Hou YP (2011) Do higher resource capture ability and utilization efficiency facilitate the successful invasion of native plants? *Biological Invasions* 13, 869–881.
- Schumacher E, Kueffer C, Edwards P, Dietz H (2009) Influence of light and nutrient conditions on seedling growth of native and invasive trees in the Seychelles. *Biological Invasions* **11**, 1941–1954.
- Souza-Alonso P, Guisande-Collazo A, González L (2015) Gradualism in Acacia dealbata Link invasion: Impact on soil chemistry and microbial community over a chronological sequence. Soil Biology and Biochemistry 80, 315–323.
- Stark JM, Norton JM (2014) The invasive annual cheatgrass increases nitrogen availability in 24-year-old replicated field plots. *Oecologia* **177**, 799–809.
- Sterner RW, Elser JJ (2002) *Ecological stoichiometry: the biology of elements from molecules to the biosphere*. (Princeton, NJ, USA: Princeton University Press).
- Van Kleunen M, Weber E, Fischer M (2010) A meta-analysis of trait differences between invasive and non-invasive plant species. *Ecology Letters* **13**, 235–245.
- Vilà M, Espinar JL, Hejda M, et al. (2011) Ecological impacts of invasive alien plants: a meta-analysis of their effects on species, communities and ecosystems. *Ecology Letters* **14**, 702–708.
- Vitousek PM, Porder S, Houlton BZ, Chadwick OA (2010) Terrestrial phosphorus limitation: Mechanisms, implications, and nitrogen-phosphorus interactions. *Ecological Applications* **20**, 5–15.
- Warton DI, Wright IJ, Falster DS, Westoby M (2006) Bivariate line-fitting methods for allometry. *Biological Reviews* **81**, 259–291.
- Yuan ZY, Chen HYH (2015) Decoupling of nitrogen and phosphorus in terrestrial plants associated with global changes. *Nature Climate Change* **5**, 465–469.
- Zabinski CA, Quinn L, Callaway RM (2002) Phosphorus uptake, not carbon transfer, explains arbuscular mycorrhizal enhancement of *Centaurea maculosa* in the presence of native grassland species. *Functional Ecology* **16**, 758–765.
- Zenni RD, Hoban SM (2015) Loci under selection during multiple range expansions of an invasive plant are mostly population specific, but patterns are associated with climate. *Molecular Ecology* **24**, 3360–3370.

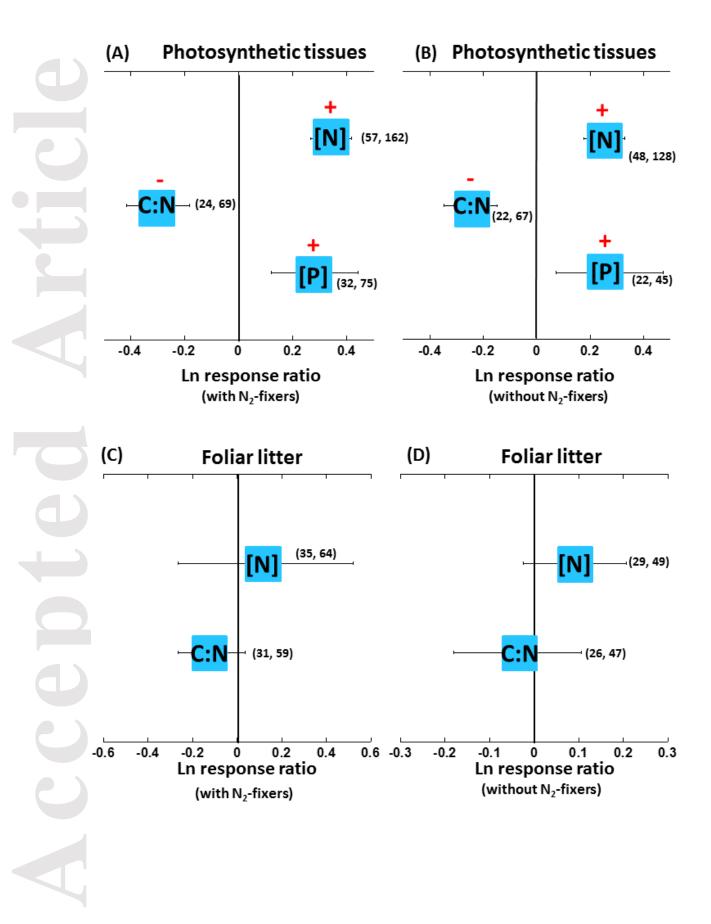
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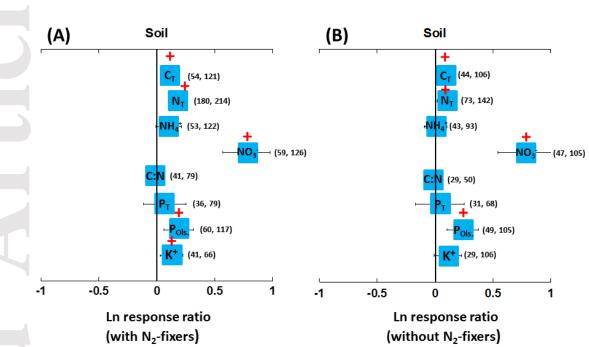
Figure 1. Ln response ratios of N and P concentrations and the C:N ratio in photosynthetic tissues (including data for N_2 -fixing plants) (A), and excluding the data for N_2 -fixing plants (B). Ln responses ratios of N concentrations and the C:N ratio in foliar litter to plant invasion for the entire data set (including data for N_2 -fixing plants) (C) and excluding the data for N_2 -fixing plants (D) to plant invasion. Values are means and 95% confidence intervals. Plus (+) and minus (–) signs represent positive and negative log response ratios, respectively, when the corresponding ln response ratio confidence intervals do not overlap with zero value. Zero in the X-axes represents neutral response ratio that means equal values in native than in invasive species. The numbers between brackets indicate the number of articles and studies (each article can have more than one single study), respectively, used in the meta-analysis of each variable.

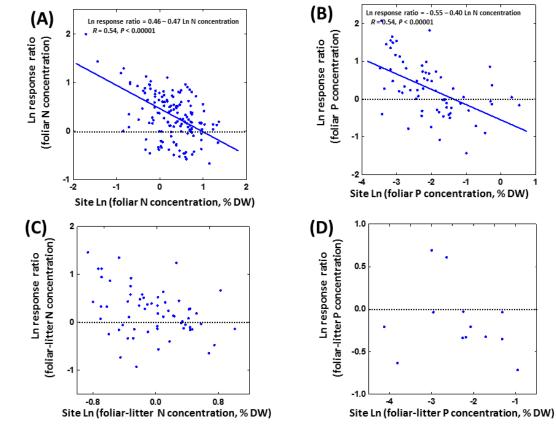
Figure 2. Ln response ratios of soil concentrations of extractable potassium (K^+), ammonium (NH_4^+), nitrate (NO_3^-), P-Olsen, and total P, N and C and the soil C:N ratio to plant invasion for the entire data set (including data for N_2 -fixing plants) (A) and excluding the data for N_2 -fixing plants (B). Values are means and 95% confidence intervals. Plus (+) and minus (–) signs represent positive and negative log response ratios, respectively, when the corresponding ln response ratio confidence intervals do not overlap with zero value. Zero in the X-axes represents neutral response ratio that means equal values in native than in invasive species. The numbers between brackets indicate the number of articles and studies, respectively, used in the meta-analysis of each variable.

Figure 3. Relationships between In response ratio of foliar N and P concentrations and the total N (A) and P (B) concentrations in the leaves of native plants, and relationships between the In response ratio of the foliar-litter N and P concentrations and the total N (C) and P (D) concentrations in the leaf litter-tissues of native plants based on percent dry wright (%DW). Dotted line highlights the zero value of In response ratio (equal values of the corresponding variable for native and in invasive species or for soils under them).

Figure 4. Relationships between the In response ratio of soil NO_3^- and the site soil NO_3^- concentration (A), between the In response ratio of soil total N and site soil total N concentration (B), between In response ratio of soil P-Olsen and site soil P-Olsen concentration (C) and between In response ratio of soil K⁺ concentration and site soil K⁺ concentration (D). Dotted line highlights the zero value of In response ratio (equal values of the corresponding variable for native and in invasive species or for soils under them.







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