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- 1 Thresholds in decoupled soil-plant elements under changing climatic conditions
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Abstract

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Background and aims Aridity has increased in the past decades and will probably 15 continue to increase in arid and semiarid regions. To decipher plant and soil capacity 16 to retain metal cations when climate evolves to more arid conditions. 17 Methods We analyzed K, Na, Ca, Mg, Fe, Mn, Zn and Cu concentrations in 580 soil 18 samples and 666 plant (shoot and root) samples along a 3600 km aridity gradient in 19 20 northern China. Results The concentrations of soil exchangeable K, Mg, Mn, Fe and Cu clearly 21 22 decreased with increasing aridity due to the relationships of aridity with soil clay content and soil pH. Increases in exchangeable Na and Ca concentrations at mid- and 23 high-aridity levels are probably due to the soil salinization, whereas increased 24 25 exchangeable Fe concentrations at extreme aridity level may be more related to a reduced pH at very high aridity. Element concentrations in both plant shoots and roots 26 were unrelated to soil exchangeable element concentrations; instead they increased 27 monotonously with increasing aridity, corresponding with decreases in plant size and 28 shoot/root ratios. The shoot/root mineralomasses ratios in general increased with 29 increasing aridity. The proportional higher element contents in shoots than in roots 30 with increasing aridity is related to increased water uptake and/or use efficiency. 31 Conclusions The extractability of soil elements in response to changing climate varied 32 33 with the nature of specific elements and to the extent these elements are controlled by biological and geochemical processes, i.e., some decreased linearly with increasing 34 aridity, whereas others first decreased and then increased with different thresholds. 35 These contrasting effects of aridity on nutrient availability could further constrain 36 plant growth and should be incorporated into biogeochemical models. The prevailing 37 paradigm of a positive relationship between concentrations of plant and soil elements 38

- 39 needs to be reconsidered under changing climatic conditions.
- 40 **Keywords** Aridity; Biogeochemical cycles; Clay; Climate change; Soil pH;
- Threshold.

Introduction

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44 Aridity has increased in the past decades and will probably continue to increase in arid and semiarid regions (Dai 2013). Such climatic changes have considerable 45 influences on global biogeochemical cycles and ecosystem development (Sardans and 46 Peñuelas 2007). Studies of the variations of concentrations of elements in the soil-47 plant system in relation to changes in aridity could enhance our ability to understand 48 and predict how ecological processes and biota will respond to global climate change 49 (Han et al. 2011; Vicente-Serrano et al. 2012; Zhang et al. 2012). To date such 50 knowledge is largely limited to essential elements such as nitrogen (N), phosphorus 51 52 (P) and sulfur (S) (Duval et al. 2013; Luo et al. 2016; Peñuelas et al. 2012; Sardans et al. 2015). However, other elements, which are also important to ecosystem functions 53 and services, are rarely studied (Duval et al. 2013; van Groenigen et al. 2006). For 54 example, potassium (K) plays an important role in stomatal behavior, osmoregulation, 55 enzyme activity and cell expansion (Wang et al. 2004) and despite this it is an 56 understudied element in global change scenarios (Sardans and Peñuelas 2015). 57 Magnesium (Mg) and calcium (Ca), two major intracellular divalent cations, are 58 important cofactors in more than 300 enzymatic reactions such as energy metabolism 59 and protein and nucleic acid synthesis (Whitehead, 2000). Copper (Cu) is an essential 60 nutrient for plant growth and development and is a component of proteins involved in 61 electron transfer and oxygen transport (Hansch and Mendel 2009). Manganese (Mn) 62 is an essential element in plants for many functions including electron transport 63 during photosynthesis and for riboflavin, ascorbic acid and carotene formation 64 (Whitehead 2000). Zinc (Zn) is also essential for plants, e.g., for the production of 65 auxins and root development (Whitehead 2000). 66

Biogeochemical cycles of multiple elements are traditionally biologically coupled due to preservation of elemental ratios in the plants, animals and microorganisms that drive them (Elser et al. 2011; Falkowski et al. 2008; Howarth et al. 2011). However, soil carbon (C) and N cycles were found to be decoupled from P and S cycles in such situations as increasing aridity in dryland ecosystems (Delgado-Baquerizo et al. 2013; Luo et al. 2016) and changing environments (Yang et al. 2014). This is because C and N cycles are most likely driven by biological processes such as photosynthesis and biological N-fixation, whereas P and S cycles are most likely driven by physical processes because P and S are rock-derived elements (Luo et al. 2016). Other macroand micro-elements, such as K, sodium (Na), Ca, iron (Fe) and Cu, can enter terrestrial ecosystems through a number of sources including biological processes (e.g., organic matter decomposition), geochemical processes (e.g., chemical weathering, salinization or changes in soil pH) and various human activities (e.g., fertility inputs) (Austin 2011; Whitehead, 2000; Ramezanian, 2013). Yet, little attention has been paid to the responses of the cycles of other elements, especially micro-elements, to a changing environment in arid and semiarid regions. In arid and semiarid regions drought can be accompanied by increases in salinity causing the precipitation and immobilization of elements such as Fe, Mn and Zn. This effect can be even greater when salinity coincides with increases in soil pH (Elamin and Hussein, 2000). Moreover, some studies have observed that drought can be related to changes in soil pH, causing either decreases (Clark et al. 2005) or increases in microelement concentrations (Kopittke et al. 2012) depending on soil type and drought intensity and timing, highlighting other possible indirect impacts of drought on microelements and trace element plant-soil cycles.

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Increasing aridity may affect not only biogeochemical cycles but also the extent to which elements are coupled by biological processes. Increases in aridity limit the soil diffusion capacity, and thus reduce the availability of elements in soils, soil microbial activities, and plant uptake; such modifications may lead to a reduction in the concentrations of elements in plant tissues (Han et al. 2011). On the other hand, it has been widely reported that plants tend to accumulate some elements such as K⁺ and Ca²⁺ to enhance cell osmotic potential under drought stress, an important adaptation strategy for maintaining water use efficiency (Chaves et al. 2003; Xoconostle-Cazares et al. 2010). These processes illustrate how aridity can affect the biological coupling of element cycles in terrestrial ecosystems. Aridity is a fundamental driver of biotic and abiotic processes in arid and semiarid areas, and hence, variations in patterns of elements in the soil-plant system may be more sensitive to changes in aridity in these regions compared to other regions with abundant rainfall (Austin 2011; Delgado-Baquerizo et al. 2013; Schroter et al. 2005). Our previous study has showed that the plant N and P concentrations would not codevelop with soil N and P availability under changing climatic conditions (Luo et al. 2015). In the present study, to examine how elements in the soil-plant system respond to aridity, the concentrations of eight mineral elements in soils and plants were studied along a 3600 km long transect representing a considerable precipitation gradient across the arid and semiarid regions in northern China. Specifically, the present study aims to demonstrate spatial patterns of metal cations in relation to climatic variables and to explore the relationships between the concentrations of these elements in plants and in top soils.

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Material and methods

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Transect and site description

In early August 2012, a field sampling campaign was conducted in 58 locations with a mean interval of 65 km between them along a 3600 km transect (West-East) in northern China (Fig. 1). The topography of the study area consists of gently rolling hills and tablelands, with elevations varying from 1500 m in the west to 700 m above sea level in the east. The sampling sites were randomly selected with an interval of 50-100 km. The geographic ranges for the data set are 39.9° N to 50.1° N (latitude) and 90.5° E to 120.4° E (longitude) (Fig. 1). Over this area, the climate is predominantly continental arid and semiarid; the annual potential evapotranspiration (PET) varies from 1229 mm (West) to 751 mm (East) and the mean annual precipitation (MAP) increases from 38 mm (West) to 436 mm (East) (see Fig. S1). The main vegetation types were shrublands, desert grasslands, typical steppe grasslands and meadow grasslands progressing from West to East. The plant species richness increased from west to east ranging from 5 to >25 species per m². The dominant plant species were *Reaumuria* spp. and *Salsola* spp. in the western region and Stipa spp., Aropyron spp., and Cleistogenes spp. in the eastern region. Soil types were predominantly arid, sandy, brown *loesials* rich in Ca, and belong to the chestnut brown and gray-brown desert soil group. Sampling sites were selected 500-1000 m away from major roads and human habitation, subjected to minimal grazing or other anthropogenic disturbances. The latitude, longitude and elevation for each sampling site were recorded with a GPS device (eTrex Venture, Garmin, USA). For more details on the sampling sites were refer to Luo et al. (2013) and Wang et al. (2014)

Sampling

At each site, two parallel 50 m × 50 m plots with a distance of approximately 1 km were selected and five 1 m × 1 m subplots were set within each plot (see Fig. 1). The five subplots were located at the four corners and in the center of the plot. In each subplot, after removing surface litter, soil samples (to 10 cm depth) were collected from ten randomly selected locations, using a soil core (2.5 cm diameter). We therefore collected a total of 580 bulk soil samples from 58 sampling sites along the large-scale transect. Soil samples were homogenized by hand mixing and then separated into two subsamples: one was stored in a plastic bag in a refrigerator at 4 °C for incubation experiments; the other was stored in a cloth bag at room temperature for soil chemical analyses.

In each subplot, the maximum height (cm) of plants belonging to each of the three dominant genera, *Stipa*, *Cleistogenes*, and *Agropyron*, was measured with a ruler. Then five to ten mature and healthy individuals of each genus were selected and extracted by pushing a soil cylinder (25 cm in diameter and 30 cm in depth) into the soil surrounding an individual plant sample and digging the plant out with a spade (Luo et al. 2013; 2015). Above- and below-ground tissues were carefully separated and then stored in paper bags separately. We collected a total of 666 plant samples (shoot and root) across the aridity gradient. The aboveground parts of the three genera in each plot were harvested and separated to calculate the aboveground biomass of each genus (g m⁻²). Within the same day, plant material sampled was dried at 105 °C for 30 min in order to minimize respirations and decomposition, and then stored at 4 °C until further processing and analyses in the laboratory. More details can be found in Luo et al. (2013) and Wang et al. (2014).

Measurements

Soils were passed through a 2 mm sieve, and fine roots and plant debris were removed. Microbial biomass C (MBC) and N (MBN) contents were analyzed with the fumigation-extraction method (Vance *et al.*, 1987). An aliquot of 10 g fresh soils (<2.0

mm) were used to measure soil pH in water (1:2.5 soil to solution).

The exchangeable K, Na, Ca and Mg concentrations were measured by extracting air-dried soils (2.5 g; <2.0 mm) with 50 ml NH₄OAc (1 M; pH=7.0), and the exchangeable Fe, Mn, Cu and Zn concentrations were measured by extracting air-dried soils (10 g; <2.0 mm) with 20 ml diethylene triamine penlaacetic acid (DTPA; pH=7.3). Concentrations of all eight elements in the soil extracts were measured by atomic absorption spectrometry (AA6800, Shimadzu, Japan). Soil inorganic C (SIC) concentrations were measured by measuring the volume of CO₂ released from air-dried soil (10 g; <2.0 mm) after treatment with 8 ml HCl (2 M) at room temperature. The SIC content was used to represent the carbonate content in our study. Soil available sulfur (SAS) concentrations were measured by extracting air-dried soils (10 g; <2.0 mm) with 50 ml CaCl₂ (0.15 %) and the contents in the soil extracts were determined using the turbidimetric method.

Soils (10 g) were fractionated into sand (particle size, 50-300 mm), silt (2-50 mm) and clay (<2.0 mm) using the ultrasonic energy method (Roscoe *et al.*, 2000). Clay content was expressed as a weight percentage of the oven-dried soil.

Soils were air-dried and ground to pass through a 1 mm sieve. Then SOC and STN concentrations were analyzed using an elemental analyzer (2400II CHN elemental analyzer; Perkin-Elmer, USA) at the Stable Isotope Facility of the University of California, Davis after removing carbonates using HCl (0.5 M).

Plant shoot and root samples were washed with deionized water and dried at 65 °C

to constant weight, and then ratios of shoot to root were measured as the dry root biomass (g) divided by the dry shoot biomass (g). Root samples were cleaned of excess soil by sonicating 3-5 g roots in 15 ml centrifuge tubes for 30 minutes in ultrapure (18 MV) water. The washed roots were again oven-dried at 65 °C to a constant weight. All plant materials were then ground and passed through a 1 mm sieve for measurement of elements.

Dried soil samples (100-150 mg; <1.0 mm) and plant samples (150-200 mg; <1.0 mm) were both acid digested with a mixture of acids (HNO₃, HClO₄ and HF, in a proportion of 5:1:2 (v/v/v) for soil samples and 5:1:0 for plant samples) in a microwave oven. Microwave digestion was performed until the sample was dissolved into the solution. The concentrations of K, Na, Ca, Mg, Fe, Mn, Zn and Cu were then measured either using inductively coupled plasma mass spectrometry (Perkin Elmer, ELAN-6000) or inductively coupled plasma emission spectroscopy (Perkin Elmer, OPTIMA 3000 DV).

Climate data

MAP and PET were extracted from a global climate dataset from http://www.worldclim.org/. Aridity was defined as 1-AI, where AI, the ratio of precipitation to potential evapotranspiration, is the aridity index (Delgado-Baquerizo et al. 2013).

Statistical analysis

Before numerical and statistical analysis, all variables (K, Na, Ca, Mg, Fe, Mn, Zn and Cu concentrations in soils and plants) were averaged at the site level. Some of these dataset were log₁₀-transformed to meet distributional assumptions underlying the statistical modeling. To demonstrate the effects of soil parent materials on the

patterns of soil exchangeable elements, ordinary least squares (OLS) linear regressions were explored between soil total elemental concentrations and the corresponding exchangeable concentrations along the transect. In addition, linear regressions were used to explore relationships between aridity and the ratios between soil exchangeable- and total elemental concentrations. Linear or curvilinear (quadratic) regressions were used to relate each soil exchangeable element to aridity to explore the effects of climate regimes on patterns of soil exchangeable elements in the present study. We found that the relationships between soil exchangeable concentrations of K, Mg, Mn, Zn and Cu and aridity were well described by linear regressions. Then OLS regressions were used to relate exchangeable concentrations of these five elements to soil pH value and contents of SIC, clay, SOC, STN, MBC and MBN. We found that the relationships between soil exchangeable Ca, Na and Fe concentrations and aridity were well described by a second-order polynomial, with thresholds at aridity being 0.65 for Ca, 0.63 for Na and 0.83 for Fe. Therefore, OLS linear regressions were used to relate soil exchangeable Ca, Na and Fe concentrations to soil pH value and contents of SIC, clay, SOC, STN, MBC and MBN above and below their thresholds. We further explored the relationships of aridity with soil pH value, and with concentrations of SIC, SAS and STN using linear regressions. To show the effects of climatic variables on plant element patterns, OLS linear regressions were used to relate aridity to the contents of plant elements in shoots and roots for the three genera. Maximum plant height and plant shoot/root ratios in relation to aridity were also explored with linear regressions. Total site mineralomasses were defined as the total contents of the eight studied elements (K, Na, Ca, Mg, Fe, Mn, Zn and Cu) in the biomass of each site by adding the contents in

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the plants of the three main genera per plot. Linear regressions were used to relate the 234 total site mineralomasses and shoot/root mineralomasses ratios with aridity. 235 All statistical univariate analyses were carried out with SPSS11.0 (SPSS, Inc., 236 USA, 2001). 237 The aridity, soil clay content and soil pH were analyzed as factors explaining the 238 maximum variability of soil exchangeable concentrations and soil exchangeable/total 239 soil concentrations ratio of K, Na, Ca, Mg, Fe, Mn, Zn and Cu by structural equation 240 modeling (SEM). This analysis provided information for the direct, indirect and total 241 242 effects of the exogenous variables on the endogenous variables. We fitted the different models using the sem R package and determined the minimum adequate model using 243 the Akaike information criterion. Standard errors and the significance level (P value) 244 of the total, direct and indirect effects were calculated using bootstrapping (1200 245 repetitions). 246

Results

Soil and plant elements in relation to environmental variables						
249	There were no significant relationships between total and exchangeable soil element					
250	concentrations (P >0.05, Fig. S3) except for Ca (R ² =0.72; P <0.001, Fig. S3B) and Cu					
251	(R^2 =0.12; P <0.01, Fig. S3H). The ratios of soil exchangeable to total elements in					
252	surface soils decreased from wetter to intermediate aridity sites; thereafter, K, Na and					
253	Fe increased whereas the other elements continued to decline (Fig. S4).					
254	The concentrations of soil exchangeable K, Mg, Mn and Cu decreased with					
255	increasing aridity (all <i>P</i> <0.01, Fig. 2). The concentrations of these four elements were					
256	positively correlated with MBC, MBN, SOC, STN, and soil clay contents, but					
257	negatively correlated with soil pH and SIC (Table 1).					
258	The concentrations of soil exchangeable Ca, Na and Fe showed a concave-shaped					
259	trend with aridity, i.e., they first decreased and then increased with aridity thresholds					
260	of 0.65 for Ca, 0.63 for Na and 0.83 for Fe (all P <0.05, Fig. 2). When the aridity was					
261	lower than these thresholds, Ca, Na and Fe concentrations were all found to be					
262	positively correlated with MBC, MBN, SOC, STN and soil clay (all <i>P</i> <0.05, Table 1)					
263	Exchangeable Ca and Na were also positively correlated with SIC when the aridity					
264	was above their thresholds (P <0.05, Table 1). When aridity was higher than their					
265	thresholds, Ca, Na and Fe concentrations were all negatively correlated with MBC					
266	and MBN concentrations, SOC and STN concentrations and soil clay (P<0.05, Table					
267	1). Fe concentrations were negatively correlated with soil pH and SIC below the					
268	threshold aridity of 0.83 (both <i>P</i> <0.05, Table 1).					
269	Unlike the patterns of exchangeable elements in soils, the concentrations of the					
270	eight elements in both plant shoots and roots increased consistently with increasing					

aridity along the aridity transect (Fig. 3).

When the total contents of the eight studied elements in the biomass of each site were analyzed, we observed that the total site mineralomasses of K, Na, Zn and Cu decreased with aridity (Fig. S5). Thus, as a general rule and notwithstanding the high concentrations in shoots and roots, with increasing aridity the total contents in stand biomass tend to decrease. Interestingly, the shoot/root ratio of the total site mineralomasses did not change with aridity (data not shown). When these correlations were studied at the genus level, the patterns of mineralomass with aridity were different than those observed with whole stand biomass. Aridity was not related with total content of any of the eight studied elements in any of the three dominant genera throughout the sites, but aridity was positively related with Na, Fe, Mn and Zn shoot/root mineralomass ratios in *Stipa* spp, with Ca, Na, Mg and Mn shoot/root mineralomass ratios of *Cleistogenes* spp., and with Mn shoot/root mineralomass ratios of *Agropyron* spp. (Fig. S6).

Multivariate soil analyses

The SEM showed that aridity had indirect significant relationships with soil exchangeable concentrations of K, Na, Ca, Fe, Mn and Cu by way of its negative relationship with soil clay contents and positive relationship with soil pH (Fig. 4 and 5). Aridity had also a significant positive relationship (direct, not related to previously commented indirect relationships) on soil Na and Ca. Thus, aridity affected soil exchangeable concentrations of the studied elements mainly due to its relationships with soil clay content and soil pH. The total linear relationship of aridity with soil exchangeable K, Mg, Mn, Fe, Cu and Zn concentrations was negative, and was positive with soil exchangeable Na and Ca concentrations (Fig. 4 and 5).

The SEM showed that exchangeable/total concentration ratios were indirectly affected by aridity through its own indirect relationships with soil clay content and soil pH (Fig. 6 and 7). Aridity had also a direct effect on concentrations not related with those indirect effects. The total relationships of aridity with ratios were negative for all elements except for Na, which had a positive relationship (Fig. 6 and 7).

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Discussion

Effects of element pool on element availability

Soil available concentrations of mineral nutrients are almost entirely derived from parent materials in terrestrial ecosystems (Foulds 1993). Hence, soil total element pools play a critical role in availability. However, we found that most soil exchangeable element patterns were unrelated to total element patterns along the climatic gradient (Fig. S3), implying that the exchangeable fractions were not directly correlated with element pool size in the parent materials. The lack of relationships of soil total and exchangeable element contents may be attributed to variations in weathering rates from parent material (and hydrologic controls on such rates) across climate and soil gradients, which might have masked the effects of soil parent materials on element availability. In arid areas succession can be viewed as a process through which biota accumulate enough nutrients (West, 1981). This would be related to the drop in mineral weathering with aridity rise (West, 1981). The lack of weathering as a result of the short wet periods complements the salt neo-formation during the intense and long dry periods (Verhge, 2009). Consistently, previous studies have shown that soil element availability in dry ecosystems was also determined by weathering rates, which releases exchangeable forms of elements from the

lithosphere, as opposed to being determined by the pool size of elements in the parent materials (Schlesinger and Bernhardt 2013). A more detailed information on the parent material (geology) and its depth is warranted for a better understanding of the mechanism underlying the absence of relationship between soil total and extractable mineral content (Vitousek and Chadwick 2013). The mineralization of organic matter is other source of available bio-elements. In arid and semi-arid environments the mineralization rates of organic matter decays by several processes such a decrease in soil microbial activity and soil enzyme activity (Sardans & Peñuelas, 2005, 2013), which is frequently associated to the production of more recalcitrant litter and soil accumulation of some bio-elements (Sardans & Peñuelas, 2007; 2013; Sardans et al., 2008). However, there was no information regarding these factors along the largescale soil transect in our study. Climatic controls on element availability Soil exchangeable K, Mg, Mn and Cu concentrations were found to be negatively correlated with aridity across the climatic gradient (0.4<aridity<1), whereas Ca, Na and Fe concentrations were only negatively correlated with aridity in regions when the aridity was below a certain level (i.e., aridity=0.65 for Ca, 0.62 for Na and 0.83 for Fe) (Fig. 2). There are various controlling mechanisms that could explain these patterns. Firstly, biological mineralization releasing exchangeable element forms from organic matter is generally considered as a major limiting factor determining exchangeable element contents (Schlesinger and Bernhardt 2013). Hence, patterns of soil exchangeable elements mainly depend on soil biological mineralization rates associated with microbial activity. This hypothesis was partly supported in our study by the positive relationships between soil exchangeable elements and MBC and MBN along the climatic gradient (Table 1). Moreover, our results also showed that when aridity rises SOC decreases with aridity. Therefore the lower rates of mineralization

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with higher aridity could be related to less organic carbon in soil to be mineralized and/or less microbial biomass to produce enzymes and to mineralize. Secondly, biological cycling shaping the vertical distributions of rock-derived elements can move micronutrients upwards because most rock-derived elements are taken up by plant roots, transported into aboveground biomass and recycled to the soil surface through litterfall (Jobbágy and Jackson 2004). Biological cycling processes decrease with increasing aridity (Moyano et al. 2013, Wang et al. 2014), which can to some extent explain the decreased rock-derived minerals with decreasing precipitation in our study. Thirdly, soil organic matter complexation is considered to be a major process in the preservation of soil elements in most temperate soils (Oades 1988). Soil organic matter can absorb many soil minerals and protect them from being lost. Hence, the declines in exchangeable element concentrations in the surface soils with increasing aridity were partly associated with decreasing organic matter. The importance of soil organic matter in this regard can be further evidenced by the positive relationships between the concentrations of most exchangeable elements and soil organic matter content along the transect in our study (Table 1). Fourthly, soil clay content also plays an important role in the retention of soil elements at the soil surface (Tiller et al. 1984). We found that the concentrations of most soil elements increased with increased soil clay content, the latter of which was inversely related with aridity. Lastly, soil pH has an effect on the solubility or retention of minerals in soils, with a greater retention rate and lower solubility of metal cations occurring at high soil pH (Martinez and Motto 2000). SEM analyses showed that drought had indirect, negative relationships with soil availability of the eight studied elements by way of its negative relationships with soil clay content and its positive relationship with soil pH (Table 1). Thus, taken together, the combined effects of these

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environmental variables result in an ultimate decline of the availability of the considered soil elements with increasing aridity along the transect. Moreover, as commented previously, previous studies have shown that soil element availability decreases with aridity because of the decreases in minerals weathering (West, 1981; Schlesinger and Bernhardt 2013).

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Our results showed that soil exchangeable concentrations of Ca, Na and Fe increased in the extreme range of aridity (mainly Ca and Na) (Fig. 2) and decoupled from K, Mg, Mn and Cu concentrations (opposite trend) when the aridity was higher than a certain threshold (i.e., aridity=0.65 for Ca, 0.62 for Na and 0.83 for Fe) (Fig. 2). We did not find evidence for a clear positive relationship between soil exchangeable concentrations of Ca, Na and Fe and biological variables such as MBC, MBN, SOC, and STN (Table 1), which indicated that patterns of soil exchangeable Ca, Na and Fe shifted from biologically-controlled factors (microbial activities and soil fertility) to geochemically- and geophysically-controlled factors (evaporation and carbonate equilibrium) with increasing aridity. Relatively high aridity can generally promote soil drying and alter hydrological transport processes, increasing salinity. Calcium (Ca) and Na in particular as relatively mobile and abundant elements in soils can be transported from soil depths and redistributed to the top soil layers as a result of hydraulic redistribution. Higher evapotranspiration can transport minerals passively to the aboveground tissues, from which minerals will be deposited to the soils. In addition, increases in aridity can produce large areas of bare soil, enhancing wind erosion and physical weathering in drylands. In support of this explanation, concentrations of soil SO₄²⁻ and CO₃²⁻ were found to accumulate under such dry conditions along our transect (see Fig. S7). In dry conditions Na and Ca accumulate in the soil surface forming salt crystals and carbonates, which in turn contribute to the

production of rock debris and thereby contributing to the release of more rock-derived elements (Rodriguez-Navarro and Doehne 1999). A decrease in soil pH at extremely dry sites (at aridity above 0.78) (see Fig. S8) can increase the solubility of Fe through pH buffering by equilibria of H₂CO₃, HCO₃⁻, and CO₃²- (Bloom, 2000). This may be one plausible reason for abruptly higher Fe concentrations at the driest sites of the present study. Moreover, dry climatic conditions favor the formation and accumulation of these compounds due to low effective precipitation, since precipitation can dissolve and leach these compounds from the soil profile.

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Soil properties and processes may change suddenly and/or nonlinearly in response to extrinsic differences in environmental forcing such as precipitation and temperature; such responses have been characterized as "pedogenic thresholds" in a variety of complex ecosystems (Vitousek and Chadwick 2013). Our results showed that Na and Ca in soil extracts decreased to an intermediate aridity and Fe concentrations in soil extracts decreased to the end of the studied range of aridity and then increased at greater aridity, forming thresholds (i.e., aridity=0.62 for Na, 0.65 for Ca and 0.83 for Fe) marking a minimum of a concave curve. Similarly, along a climatic gradient in the Sierra Nevada Range, Dahlgren et al. (1997) found that primary mineral weathering and clay mineral formation increased abruptly within a relatively narrow climatic zone due to a favorable combination of temperature and precipitation. Chadwick et al. (2003) analyzed soil properties and processes along an arid to humid climosequence on Kohala Mountain, Hawaii and identified pedogenic thresholds where mineral weathering and soil properties changed greatly with increased rainfall. Vitousek and Chadwick (2013) demonstrated that concentrations of silicon (Si), aluminum (Al), and Fe in surface soils along a rainfall gradient in Hawaii declined with increasing precipitation from 260 mm yr⁻¹ to approximately 1700 mm

yr⁻¹, thereafter, Fe concentrations were enhanced abruptly whereas Si and Al concentrations continued to decrease.

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Minor environmental forcing at pedogenic thresholds could have profound and long-term consequences on ecosystem functions (Lenton et al. 2008; Scheffer et al. 2009; Scheffer et al. 2012). Identifying changes in the relationships between aridity and ecosystem element cycles can reveal critical vulnerability of arid and semiarid ecosystems to global climate change (Scheffer et al. 2009). Due to the profound influence of elements on plant growth and maintenance and reproduction of terrestrial ecosystems, the cycles of multiple elements are at the core of ecosystem functions. The present study has indicated that a small climate change can affect ecosystem processes by differently impacting on distinct elements and this can be even greater at specific aridity thresholds (aridity=0.62 for Na; 0.65 for Ca and 0.83 for Fe); once local aridity passes beyond these turning points, the decoupling among different elements can be larger. This would force into a long process of recovery (Scheffer et al. 2009). Based on our results, a conceptual model was proposed to show the effects of environmental factors on soil exchangeable element concentrations along the aridity gradient (Fig. S10). This simple model may be applied to other dryland ecosystems, but possibly with different aridity scales of change in the different elements in response to aridity and changes in the slope responses of some of the elements because of differences in soil chemical and physical properties, vegetation types, and atmospheric deposition rates across scales. Our study provides robust, direct evidence for improvement of the process-based modeling of biogeochemical cycling in arid and semi-arid areas.

The decoupling of biogeochemical cycles of multiple elements in drylands when

aridity rises and the different responses of the different element cycles in plant-soil system when aridity reaches certain levels may also have profound consequences one ecosystem structures, functions, and productivity by the stoichiometry shifts (Peñuelas and Sardans 2009). For instance, when aridity is above 0.62 (as observed in the present study) for Na this would cause a rise in the accumulation of Na in the soil surface and enhance osmotic stress. Plant would necessarily consume a lot of energy in the osmoregulation, which may cause them serious injuries, resulting in losses of many plant species and a reduction of vegetation cover (Chaves et al. 2003). These stochiometrical changes linked to the different rates of change in some elements when aridity reaches certain levels may exacerbate the negative effects of aridity on food production and the photosynthetic capacity of ecosystems increasing processes of degradation of the arid and semiarid ecosystems, with feed-back effects worsening the plant community capacity to retain nutrients, such observed in this study. Moreover, this decoupling of biogeochemical cycles with changes in aridity necessarily implies an ecosystem stoichiometry shift. For example, we have observed a positive relationship between aridity and the soil exchangeable Ca/Mg ratio ($R^2=0.26$, P<0.0001). Soil Ca/Mg ratios have been associated with variations in several cellular structures and functions in terrestrial plants (Stael et al. 2012).

Effects of climate and soil on plant elements

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In the present study, we found that plant shoot and root element concentrations increased with increasing aridity for the three genera but were not correlated with exchangeable element concentrations in soils along the transect (Fig. 3). One of the

possible reasons for this lack of relationship is the "dilution effect" of plant size (i.e., biomass) with increasing water availability (Jarrell and Beverly 1981). Higher precipitation significantly increased plant size in this water-limited ecosystem (see Fig. S9), which, in turn, could dilute element concentrations in plant tissues. Another reason for this lack of correlation between plant element and soil exchangeable element concentrations is the significantly decreased plant shoot/root ratios with increasing aridity (see Fig. S9), which may increase the ratios of plant mineral uptake to mineral demand, reducing the dependece of plant mineral content on soil mineral availability. Further, such uncoupling of soil nutrient element availability and plant nutrient content may also be attributed to the decreased vegetative cover and biomass with increased aridity (Wang et al. 2014), which may decrease resource competition pressure among plants, partly compensating for the reduced soil nutrient availability. In fact, when we analyzed total element contents in community biomass as the product between concentrations and biomasses in the species of the three dominant genera, we observed some increase of total K, Na, Zn and Mg at the beginning of the aridity gradient with a further change in the slope sign, with decreased in biomass contents of these elements. Thus, the results showed that the effects of aridity on total biomass is proportionally higher than the concentration effect under enhanced drought for these four elements. This trend was also observed for the other four elements but was insignificant. Moreover, and interestingly, the shoot/root mineralmass ratios of several studied elements (Fe, Na, Zn, Mg and Mn) increased with aridity in several species (Fig. S6). These results have at least three consequences for ecological stoichiometry. First, the fact that each plant taxon tends to allocate these elements more to shoots than roots with increasing aridity, thereby increasing favorable osmotic conditions, suggests a possible strategy for the improvement of water uptake and use

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efficiency, related with K (and other highly soluble elements such as Na) contents and concentrations (Fig. 3). Second, as this aridity-related trend affects elements differently, it generates a shift in plant organ stoichiometry. Third, changes in shoot/root allocation associated with drought can be distinct at the community and species-specific levels. In the case of this study we did not observe a shift in community shoot/root ratios along the gradient but this was observed for some elements at the species level. While species-specific increases in shoot/root ratio were related with increasing aridity, increasing aridity also diminished the presence of the *Agropyron* spp. and *Cleistogenes* spp. which had the lowest shoot/root ratios. Thus, the former effect was counteracted by the latter.

In conclusions, In general the concentrations of exchangeable elements in soils decreased with increasing aridity, but with three exceptions: Na and Ca in soil extracts decreased only up to a threshold of intermediate aridity, and Fe decreased up to a point close to the end of the studied range of aridity, but thereafter increased with greater aridity. The biogeochemical cycles of these elements, normally coupled in less arid regions, were decoupled in more arid regions at aridity values of 0.62 for Na, 0.65 for Ca and 0.83 for Fe. The decoupling appeared to be most directly associated with the balance between biological and geochemical controlling mechanisms of element cycles. Both linear and nonlinear relationships between element cycles and climate change can greatly influence the plant growth and ecology functions, especially for arid and semiarid ecosystems where climate regimes play a profound role in plant and ecosystem functions. The multiplicity of thresholds of different mineral elements may also imply that ecosystem functions such as biogeochemical processes could abruptly change at multiple points along an environmental gradient. Plant element concentrations decreased with increases in plant size and shoot/root

ratios, both associated with increasing rainfall, but concentrations had no relationships with soil element availability. Thus, in a general sense the results describe lower soil availability of elements and lower plant element stocks, resulting in a decrease in the amount of elements involved in the plant-soil cycle with increasing aridity. Depending on the element, this outcome manifested with differing intensities, and for three of the eight elements there were also interrupted or inverse patterns. The results also showed a decoupling among the studied elements with increasing aridity, which is relevant for other areas and biogeochemical and ecosystem models. Overall, our findings advance our understanding of the unique nature of cycles of multiple elements in soil-plant systems across wide gradients of environmental factors and make it possible to better parameterize complex multi-element biogeochemical models included in Earth system models.

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References

539

Austin AT (2011) Has water limited our imagination for aridland biogeochemistry? 540 541 Trends in Ecology & Evolution 26: 229-235. Bloom PR (2000) Soil pH and pH buffering. In M. Sumner (ed.) Handbook of soil 542 science. CRC Press, Boca Raton, FL. pp. B-333-B-352. 543 Chadwick OA, Gavenda RT, Kelly EF, Ziegler K, Olson CG, Elliott WC, Hendricks 544 DM (2003) The impact of climate on the biogeochemical functioning of volcanic 545 soils. Chemical Geology 202: 195-223. 546 547 Chaves MM, Maroco JP, Pereira JS (2003) Understanding plant responses to drought—from genes to the whole plant. Functional Plant Biology 30: 239-264. 548 Clark JM, Chapman PJ, Adamson JK, Lane SN (2005) Influence of drought-induced 549 acidification on the mobility of dissolved organic carbon in peat soils. Global 550 Change Biology 11: 791-809. 551 Dahlgren R, Boettinger J, Huntington G, Amundson R (1997) Soil development along 552 553 an elevational transect in the western Sierra Nevada, California. Geoderma 78: 207-236. 554 555 Dai AG (2013) Increasing drought under global warming in observations and models. Nature Climate Change 3: 52-58. 556 Delgado-Baquerizo M, Maestre FT, Gallardo A, Quero JL, Ochoa V, Garcia-Gomez 557 M, Escolar C, Garcia-Palacios P, Berdugo M, Valencia E, Gozalo B, Noumi Z, 558 Derak M, Wallenstein MD (2013) Aridity modulates N availability in arid and 559 semiarid Mediterranean grasslands. PloS One 8: e59807. 560

Duval BD, Dijkstra P, Drake BG, Johnson DW, Ketterer ME, Megonigal JP, Hungate 561 BA (2013) Element Pool Changes within a Scrub-Oak Ecosystem after 11 Years 562 of Exposure to Elevated CO₂. PloS One 8: e64386. 563 Elamin EA, Hussein AH (2000) Cooper adsorption as affected by electrolyte 564 concentration and sodium adsorption ratio in three major soil series in Sudan. An 565 Arid Zone 39: 137-143. 566 Elser JJ, Acquisti C, Kumar S (2011) Stoichiogenomics: the evolutionary ecology of 567 macromolecular elemental composition. Trends in Ecology & Evolution 26: 38-568 44. 569 Falkowski PG, Fenchel T, Delong EF (2008) The microbial engines that drive Earth's 570 biogeochemical cycles. Science 320: 1034-1039. 571 Foulds W (1993) Nutrient concentrations of foliage and soil in South-western 572 Australia. New Phytologist 125: 529-546. 573 Han WX, Fang JY, Reich PB, Ian Woodward F, Wang ZH (2011) Biogeography and 574 variability of eleven mineral elements in plant leaves across gradients of climate, 575 soil and plant functional type in China. Ecology Letters 14: 788-796. 576 Hansch R, Mendel RR (2009) Physiological functions of mineral micronutrients (Cu, 577 Zn, Mn, Fe, Ni, Mo, B, Cl). Current Opinion Plant Biology 12: 259-266. 578 Howarth R, Chan F, Conley DJ, Garnier J, Doney SC, Marino R, Billen G (2011) 579 Coupled biogeochemical cycles: eutrophication and hypoxia in temperate 580 estuaries and coastal marine ecosystems. Frontiers in Ecology and the 581 Environment 9: 18-26. 582

- Jarrell W, Beverly R (1981) The dilution effect in plant nutrition studies. Advances in
- 584 Agronomy 34: 197-224.
- Jobbágy EG, Jackson RB (2004) The uplift of soil nutrients by plants: biogeochemical
- consequences across scales. Ecology 85: 2380-2389.
- Kopittke GR, Tietema A, Verstraten JM (2012) Soil acidification occurs under
- ambient conditions but is retarded by repeated drought: Results of a field-scale
- climate manipulation experiment. Science of the Total Environment 439: 332-
- 590 342.
- Lenton TM, Held H, Kriegler E, Hall JW, Lucht W, Rahmstorf S, Schellnhuber HJ
- 592 (2008) Tipping elements in the Earth's climate system. Proceedings of the
- National Academy of Sciences of the United States of America 105: 1786-1793.
- Luo W, Elser JJ, Lü XT, Wang Z, Bai E, Yan C, Wang C, Li MH, Zimmermann NE,
- Han X, Xu Z, Li H, Wu Y, Yong J (2015) Plant nutrients do not covary with soil
- nutrients under changing climatic conditions. Global Biogeochemical Cycles 29,
- 597 doi: 10.1002/2015GB005089.
- Luo W, Jiang Y, Lü X, Wang X, Li MH, Bai E, Han X, Xu Z (2013) Patterns of plant
- biomass allocation in temperate grasslands across a 2500-km Transect in
- Northern China. PloS One 8. doi: 10.1371/journal.pone.0071749.
- Luo WT, Dijkstra FA, Bai E, Feng J, Lü XT, Wang C, Wu HH, Li MH, Han XG, Jiang
- Y (2016) A threshold reveals decoupled relationship of sulfur with carbon and
- nitrogen in soils across arid and semi-arid grasslands in northern China.
- Biogeochemistry. 127:141-153.

Martinez CE, Motto HL (2000) Solubility of lead, zinc and copper added to mineral 605 soils. Environmental Pollution 107: 153-158. 606 Moyano FE, Manzoni S, Chenu C (2013) Responses of soil heterotrophic respiration 607 to moisture availability: An exploration of processes and models. Soil Biology 608 and Biochemistry, 59: 72-85. 609 Oades J (1988) The retention of organic matter in soils. Biogeochemistry 5: 35-70. 610 Peñuelas J, Sardans J (2009) Ecology: Elementary factors. Nature 460: 803-804. 611 Peñuelas J, Sardans J, Rivas-ubach A, Janssens IA (2012) The human-induced 612 613 imbalance between C, N and P in Earth's life system. Global Change Biology 18: 3-6. 614 Ramezanian BA (2013) Influence of soil amendments and soil properties on macro-615 and micronutrient availability to microorganisms and plants. Acta Universitatis 616 Agriculturae Sueciae 30: 1652-6880. 617 Rodriguez-Navarro C, Doehne E (1999) Salt weathering: influence of evaporation 618 rate, supersaturation and crystallization pattern. Earth Surface Processes and 619 Landforms 24: 191-209. 620 Roscoe R, Buurman P, Velthorst EJ (2000) Disruption of soil aggregates by varied 621 amounts of ultrasonic energy in fractionation of organic matter of a clay Latosol: 622 carbon, nitrogen and delta C-13 distribution in particle-size fractions. European 623 Journal of Soil Science 51: 445-454. 624 Sardans J, Janssens IA, Alonso R, Veresoglou SD, Rillig MC, Sanders TG, Carnicer J, 625 Filella I, Farré-Armengol G, Peñuelas J (2015) Foliar elemental composition of 626 European forest tree species associated with evolutionary traits and present 627

628	environmental and competitive conditions. Global Ecology and Biogeography						
629	24: 240-255.						
630	Sardans J, Peñuelas J (2005) Drought decreases soil enzyme activity in a Mediterranean						
631	Quercus ilex L. forest. Soil Biology and Biochemistry 37: 455-461.						
632	Sardans J, Peñuelas J (2007) Drought changes phosphorus and potassium accumulation						
633	patterns in an evergreen Mediterranean forest. Functional Ecology 21: 191-201.						
634	Sardans J, Peñuelas J (2007) Drought changes the dynamics of trace element						
635	accumulation in a Mediterranean Quercus ilex forest. Environmental pollution						
636	147: 567-583.						
637	Sardans J, Peñuelas J (2013) Plant-soil interactions in Mediterranean forest and						
638	shrublands: impacts of climatic change. Plant and Soil 365: 1-33.						
639	Sardans J, Peñuelas J (2015) Potassium: a neglected nutrient in global change. Global						
640	Ecology and Biogeography 24: 261-275.						
641	Scheffer M, Bascompte J, Brock WA, Brovkin V, Carpenter SR, Dakos V, Held H, van						
642	Nes EH, Rietkerk M, Sugihara G (2009) Early-warning signals for critical						
643	transitions. Nature 461: 53-59.						
644	Scheffer M, Carpenter SR, Lenton TM, Bascompte J, Brock W, Dakos V, van de						
645	Koppel J, van de Leemput IA, Levin SA, van Nes EH, Pascual M, Vandermeer J						
646	(2012) Anticipating Critical Transitions. Science 338: 344-348.						
647	Schlesinger WH, Bernhardt ES (2013) Biogeochemistry: an analysis of global change.						
648	Academic press.						
649	Schroter D, Cramer W, Leemans R, Prentice IC, Araujo MB, Arnell NW, Bondeau A,						
650	Bugmann H, Carter TR, Gracia CA, de la Vega-Leinert AC, Erhard M, Ewert F,						
651	Glendining M, House JI, Kankaanpaa S, Klein RJT, Lavorel S, Lindner M,						

Metzger MJ, Meyer J, Mitchell TD, Reginster I, Rounsevell M, Sabate S, Sitch 652 S, Smith B, Smith J, Smith P, Sykes MT, Thonicke K, Thuiller W, Tuck G, 653 Zaehle S, Zierl B (2005) Ecosystem service supply and vulnerability to global 654 change in Europe. Science 310: 1333-1337. 655 Stael S, Wurzinger B, Mair A, Mehlmer N, Vothknecht UC, Teige M (2012) Plant 656 organellar calcium signaling: an emerging field. Journal of Experimental Botany 657 63: 1525-1542. 658 Tiller KG, Gerth J, Brümmer G (1984) The relative affinities of Cd, Ni and Zn for 659 different soil clay fractions and goethite. Geoderma 34: 17-35. 660 van Groenigen KJ, Six J, Hungate BA, de Graaff MA, van Breemen N, Van Kessel C 661 (2006) Element interactions limit soil carbon storage. Proceedings of the 662 National Academy of Sciences 103: 6571-6574. 663 Verheye W (2009) Soils of arid and semi-arid areas. In Land use, land cover and soil 664 sciences, Vol. VII. Verheye W (ed). ISBN: 978-1-84826-691-9. UNESCO; 67-95. 665 http://www.eolss.net/Sample-Chapters/C12/E1-05-07-16.pdf 666 Vicente-Serrano SM, Zouber A, Lasanta T, Pueyo Y (2012) Dryness is accelerating 667 degradation of vulnerable shrublands in semiarid Mediterranean environments. 668 Ecological Monographs 82: 407-428. 669 Vitousek PM, Chadwick OA (2013) Pedogenic thresholds and soil process domains in 670 basalt-derived soils. Ecosystems 16: 1379-1395. 671 Wang C, Wang X, Liu D, Wu H, Lu X, Fang Y, Cheng W, Luo W, Jiang P, Shi J, Yin 672 H, Zhou J, Han X, Bai E (2014) Aridity threshold in controlling ecosystem 673 nitrogen cycling in arid and semi-arid grasslands. Nature Communications 5, doi: 674

675	10.1038/ncomms5799.						
676	Wang S, Wan C, Wang Y, Chen H, Zhou Z, Fu H, Sosebee RE (2004) The						
677	characteristics of Na+, K+ and free proline distribution in several drought-						
678	resistant plants of the Alxa Desert, China. Journal of Arid Environments 56: 525-						
679	539.						
680	West, NE. 1981. Nutrient Cycling in Desert Ecosystems. In: Goodall, D.W., Perry, R.A.						
681	Eds. Arid-Land Ecosystems. Volume 2. pp 301-324.						
682							
683	Whitehead DC (2000) Nutrient elements in grassland. Soil-plant-animal						
684	relationships. Wallingford, UK: CABI Publishing.						
685	Xoconostle-Cazares B, Ramirez-Ortega FA, Flores-Elenes L, Ruiz-Medrano R (2010)						
686	Drought tolerance in crop plants. American Journal of Plant Physiology 5: 241-						
687	256.						
688	Yang Y, Wang G, Shen H, Yang Y, Cui H, Liu Q (2014) Dynamics of carbon and						
689	nitrogen accumulation and C: N stoichiometry in a deciduous broadleaf forest of						
690	deglaciated terrain in the eastern Tibetan Plateau. Forest Ecology and						
691	Management 312: 10-18.						
692	Zhang SB, Zhang JL, Slik JWF, Cao KF (2012) Leaf element concentrations of						
693	terrestrial plants across China are influenced by taxonomy and the environment.						
694	Global Ecology and Biogeography 21: 809-818.						
695							

Table 1 Relationships between soil exchangeable element concentrations and soil properties (soil pH, soil inorganic carbon (SIC), clay, soil organic carbon (SOC), soil total nitrogen (STN), and microbial biomass carbon (MBC) and nitrogen (MBN)) along the climatic gradient. Pearson correlation coefficients are shown.

		pН	Clay	SIC	SOC	STN	MBC	MBN
K		-0.447***	0.723***	-0.313**	0.772***	0.790***	0.674***	0.614***
Ca	Above threshold	-0.066	-0.006	0.838***	-0.560**	-0.512**	-0.562**	-0.576**
	Below threshold	0.048	0.948***	0.129	0.875***	0.916***	0.654**	0.586*
Na	Above threshold	-0.066	-0.097	0.750**	-0.458**	-0.476**	-0.515**	-0.516**
	Below threshold	-0.098	0.786**	0.145	0.634*	0.662*	0.267	0.353
Mg		-0.311*	0.650***	-0.286*	0.579***	0.590***	0.449***	0.411**
Fe	Above threshold	-0.163	-0.490**	0.231	-0.379*	-0.435*	-0.536**	-0.505**
	Below threshold	-0.645**	0.685*	-0.406**	0.835***	0.826***	0.627***	0.596**
Mn		-0.643**	0.681***	-0.417**	0.895***	0.885***	0.752***	0.700***
Zn		-0.258	0.356**	-0.051	0.406**	0.409**	0.213	0.23
Cu		-0.450***	0.654***	-0.321*	0.721***	0.699***	0.515***	0.467***

Note: 'Above threshold' represents the regions with aridity which is higher than the element specific-threshold, and 'below threshold' represents the regions with aridity which is lower than the element specific-threshold (aridity=0.65 for Ca, 0.62 for Na and 0.83 for Fe; see Fig. 2). For more details of the soil properties along the transect, refer to Fig. S9 in the supporting information section. *, P<0.05; **, P<0.01; ***, P<0.001.

Figure legends

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Fig. 1 A 3600 km transect in northern China. A total of 58 sampling points from West 707 to East were selected along the transect. Two 50 m × 50 m plots were selected and 708 five 1 m × 1 m sampling subplots were placed within each plot at the four corners and 709 the center at each site. Aridity is defined as 1-AI, where AI, the ratio of precipitation 710 to potential evapotranspiration, is the aridity index. 711 Fig. 2 Relationships between eight soil element concentrations and aridity across 58 712 713 sampling sites in northern China's arid and semiarid areas. Aridity is defined as 1-AI, where AI, the ratio of precipitation to potential evapotranspiration, is the aridity index. 714 Fig. 3 Relationships between eight plant element concentrations and aridity in 715 northern China's arid and semiarid areas. Data for the three graminoid genera sampled 716 are distinguished by color and symbol. Regression R² values are given in the Fig. 717 Aridity is defined as 1-AI, where AI, the ratio of precipitation to potential 718 evapotranspiration, is the aridity index. 719 720 Fig. 4 Diagrams of the structural equation models that best explained the maximum variance of the soil exchangeable concentrations of the eight studied elements and 721 722 aridity, soil clay content and soil pH as exogenous factors. Black and red arrows 723 indicate negative and positive relationships, respectively. Fig. 5 Total, direct and indirect effects of aridity, soil clay content and soil pH on soil 724 exchangeable concentrations of the eight studied elements. Data obtained by using the 725 bootstrap technique (with 1200 repetitions). Aridity is defined as 1-AI, where AI, the 726 727 ratio of precipitation to potential evapotranspiration, is the aridity index. 728 Fig. 6 Diagrams of the structural equation models that best explained the maximum variance of the soil exchangeable/total concentration ratios of the eight studied 729 elements and aridity, soil clay content and soil pH as exogenous factors. Black and red 730

arrows indicate negative and positive relationships respectively.

Fig. 7 Total, direct and indirect effects of aridity, soil clay content and soil pH on soil
exchangeable/total concentration ratios of the eight studied elements. Data obtained
by using the bootstrap technique (with 1200 repetitions). Aridity is defined as 1-AI,
where AI, the ratio of precipitation to potential evapotranspiration, is the aridity index.

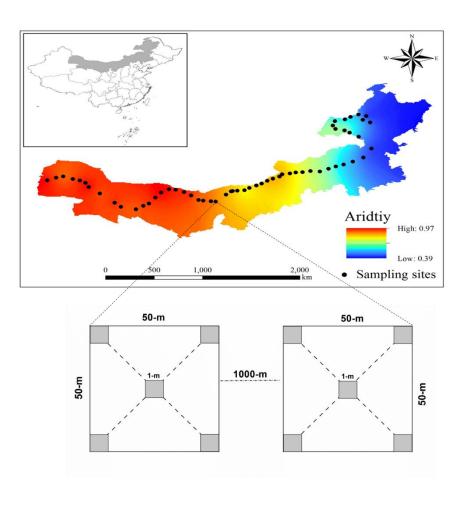


Fig. 1

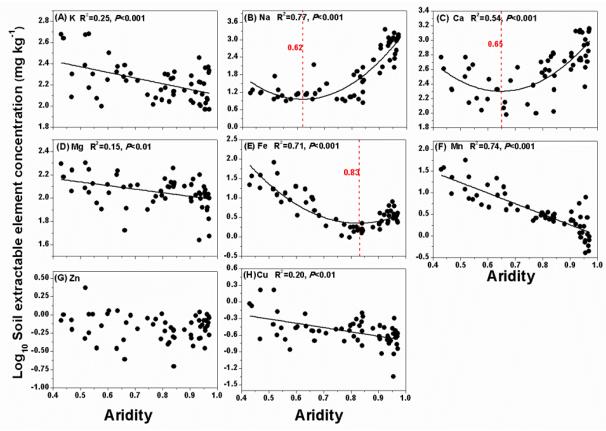


Fig. 2

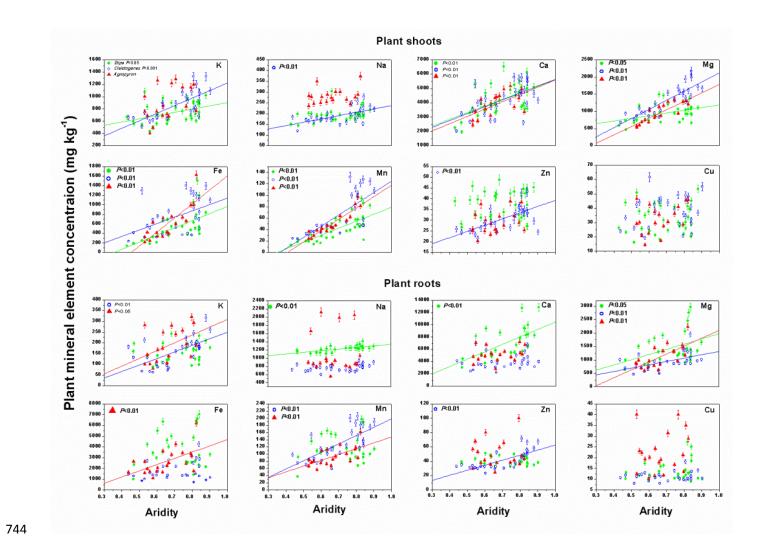
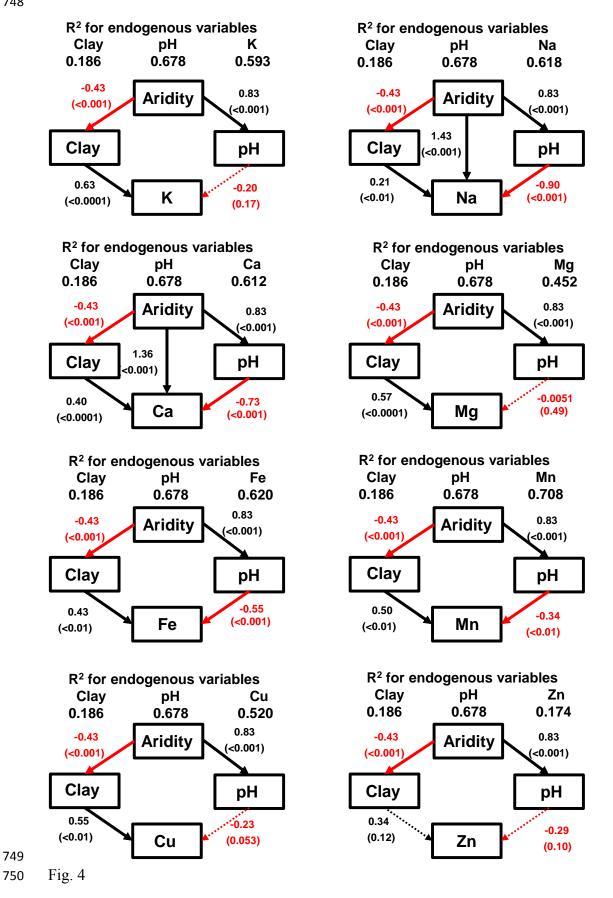


Fig. 3



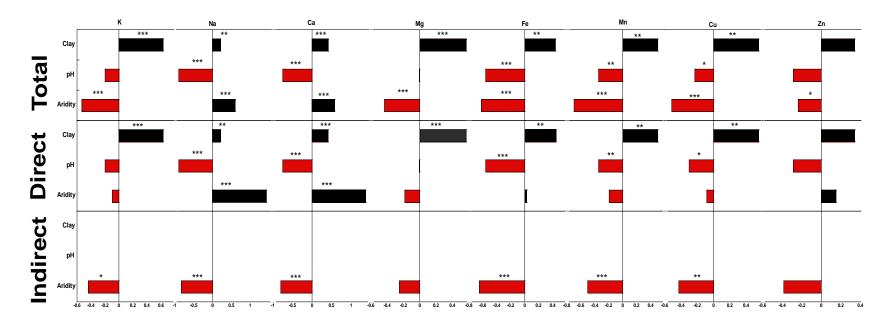


Fig. 5

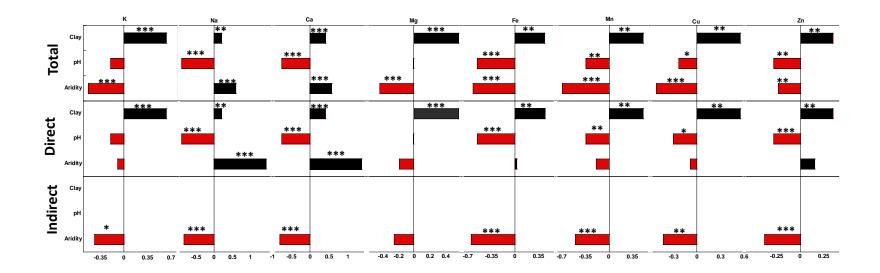
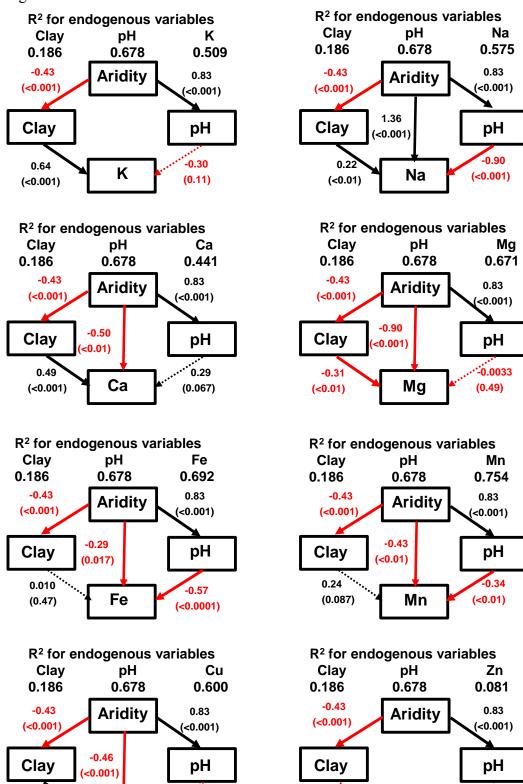




Fig. 6



0.36

(<0.01)

Cu

-0.086

(0.33)

Zn

-0.086

(0.36)

•-0.10

(0.20)

764 Fig. 7

