Understanding nutrient landscapes for giant pandas in the Qinling Mountains, China: the relationships between bamboo mineral content and giant panda habitat selection during migration

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Abstract

Understanding nutrient landscapes for giant pandas in the Qinling Mountains, China: the relationships between bamboo mineral content and giant panda habitat selection during migration. Bamboo comprises over 99% of the diet of giant pandas (Ailuropoda melanoleuca). Giant pandas face a complex nutrient landscape. They eat more than one species of bamboo and various parts of the plant, and they move seasonally to find optimal forage. Though the seasonal habitat preferences of giant pandas have long been known, the spatial and temporal nutrient gradient of bamboo between seasonal habitats remains unclear. Few studies detail the nutrient content of bamboo in relation to the seasonal habitat selection of giant pandas in the wild. In this study, we collected bamboo samples from 57 plots considering four factors (seasons, elevations, species, and plant parts). We evaluated the effect of these factors on the contents of seven bamboo mineral elements (Cu, Zn, Fe, Mn, K, Ca, and Mg) and used a non-parametric ensemble tree model to model giant pandas' presence and absence based on bamboo mineral content. Our results showed strong correlations between pairs of mineral contents (up to r = 0.69) with specific mineral elements such as Mn, consistently showing great importance in the models for differentiating the habitat selection. We also observed significant variation in mineral concentrations between seasons, bamboo species, and plant parts. Our results suggest that the studied bamboo mineral content strongly associates giant pandas' habitat preferences. Our research may be useful for the development of conservation and reserve management strategies by providing guidelines to increase giant pandas' opportunities to obtain sufficient nutrient within the Qinling region.

Key words: Giant pandas, Habitat selection, Machine learning, Mineral elements, Nutrient content, Qinling Mountains

Resumen

Comprender la distribución de nutrientes en el territorio del panda gigante en las montañas Qinling, en China: las relaciones entre el contenido de minerales del bambú y la selección del hábitat del panda gigante durante la migración. A pesar de que el bambú constituye más del 99% de la dieta del panda gigante (Ailuropoda melanoleuca), esta especie se enfrentan a un complejo patrón de disponibilidad de nutrientes, ya que consumen más de una especie de bambú y varias partes de la planta, y se desplazan de forma estacional para encontrar el alimento óptimo. Si bien las preferencias estacionales del hábitat del panda gigante se conocen desde hace tiempo, el gradiente espacial y temporal de los nutrientes del bambú entre los hábitats estacionales sigue siendo poco claro. Son pocos los estudios en los que se describe con detalle el contenido de nutrientes del bambú en relación con la selección estacional del hábitat del panda gigante en el medio silvestre. En este estudio, recogimos muestras de bambú de 57 parcelas teniendo en cuenta cuatro factores (estación del año, altura, especie y parte de la planta). Se evaluó el efecto de estos factores en el contenido de siete elementos minerales del bambú (Cu, Zn, Fe, Mn, K, Ca y Mg) y se utilizó un modelo de árbol no paramétrico para determinar la presencia y ausencia del panda gigante en función del contenido de minerales del bambú. Nuestros resultados mostraron que existen fuertes correlaciones entre pares de contenidos de minerales (hasta r = 0,69) y que algunos elementos minerales, como el Mn, son siempre un factor importante en la selección del hábitat. También observamos una variación significativa en las concentraciones de minerales entre estaciones, especies de bambú y partes de la planta. Los resultados sugieren que el contenido de minerales del bambú estudiado está fuertemente relacionado con

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las preferencias de hábitat de los pandas gigantes. Nuestra investigación puede ayudar a elaborar estrategias de conservación y gestión de reservas al ofrecer pautas que ayuden a aumentar la posibilidad de que el panda gigante pueda obtener los nutrientes que necesita en la región de Qinling.

Palabras clave: Pandas gigantes, Selección del hábitat, Aprendizaje automático, Elementos minerales, Contenido de nutrientes, Montañas Qinling

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Introduction

Bamboo comprises over 99% of the diet of wild giant pandas (Ailuroplda melanoleuca) (Sheldon, 1937; Milton, 1979; Dierenfeld et al., 1982; Tuanmu et al., 2012). Though giant pandas forage primarily on vegetation, their gut anatomy resembles that of a carnivore, with a simple stomach and short gastrointestinal tract which greatly limits its ability to digest fiber. The poor nutrition content of bamboo, which is low in protein and high in fiber, requires giant pandas to consume large quantities of bamboo to meet their dietary requirements (Milton, 1979; Dierenfeld et al., 1982; Tuanmu et al., 2012). Climate change and shifts in the phenology and physiological conditions of bamboo may exacerbate the challenges of finding suitable habitat to meet giant pandas dietary requirements (Liu et al., 1999; Li and Manfred, 2001; Hunter et al., 2003; Songer et al., 2012; Hull et al., 2014).

Given the dependence of giant pandas on bamboo, a deeper understanding of the nutrient landscape faced by giant pandas is critical to conservation efforts (Mainka et al., 1989; Reid et al., 1989; Reid and Hu, 1991; Taylor et al., 1991). Previous studies highlighted a variety of factors (seasons, elevations, species, and bamboo plant parts) affecting the mineral content of bamboo (Fu et al., 1990; Liu, 2008; Sun et al., 2010; Wang et al., 2010, 2013). Most studies, however, focused on the effect of only one or two of these factors and presented only the empirical measurements of the chemical composition of bamboo (Taylor and Qin, 1993; Li et al., 2007; Wang et al., 2009; Wu et al., 2009). A comprehensive analysis of the relationship between multiple covariables and bamboo nutrients is needed (Liu, 2001; Liu et al., 2002, 2005; Finley et al., 2011; Hull et al., 2011).

Giant panda's habitat selection in the Qinling Mountains has been extensively documented. Giant pandas migrate in late spring from their home range at low elevations to their summer home range at higher elevations and return to their low elevation range in autumn (Schaller et al., 1989; Liu et al., 2002, Lu et al., 2007; Qi et al., 2011, 2012). It has been hypothesized that this migration pattern evolved to facilitate access to either abundant or more nutritious forage (Zhang et al., 2006, 2014; Sims et al., 2007; Viña et al., 2010; Wang et al., 2010). However, little work has been done to understand the relationships among the nutrient composition of bamboo and giant pandas habitat preferences.

In our study, we measured the nutrient content of bamboo across seasons and elevation gradients. We collected samples from three elevation strata in spring, summer and autumn. Each season, giant pandas are present in one of the strata and absent in the other two. This enabled us to examine the nutrient landscape of giant pandas at various locations and seasons and to explore the statistical association between bamboo nutrient content and giant pandas' habitat choices. Furthermore, it allowed us to compare the relative contribution of individual mineral elements to giant pandas' habitat selection.

We first tested the correlation between seven mineral elements. We then examined the effect of the four factors (season, elevation, species, and plant

part) on the mean concentrations of each mineral. Finally, we modeled the presence and absence of giant pandas using the mineral element contents with a non–parametric ensemble tree model. For each season we built one model and evaluated the importance of predictive variables to the model. We sought to answer the following research questions: (1) How do each of the seven mineral elements change with season, elevation, part, and species? (2) What are the relative associations between the seven mineral elements across landscapes and giant pandas' seasonal selection of habitat?

Material and methods

Foping Nature Reserve (FNR) in Shaanxi Province was founded in 1978 in the center of a giant panda reserve network in the Qinling Mountain range. FNR (32°32' - 33°43' N, 107°41' - 107°56' E) covers an area of 293 km² and is one of earliest reserves established for giant pandas. The annual temperature at FNR was 11.5 °C (mean min = -3 °C in January, mean max = 28 °C in July) with elevation ranging between 1000 m and 2900 m. FNR is home to three bamboo species: arrow bamboo (Fargesia qinlingensis) dominating at higher elevation (1,700-2,900 m), wooden bamboo (Bashania fargesii) mostly found at lower elevation (1,000-1,900 m), and dragon-head bamboo (Fargesia racocephala) restricted to a small area in the southeast corner of the FNR (1,000-1,800 m). Arrow bamboo is the giant pandas' only food resource in summer, while they forage primarily on wooden bamboo and dragon-head bamboo in other seasons (State Forestry Administration of the People's Republic of China, 2006; Zhang et al., 2014).

Guanyinshan Nature Reserve (GNR) was founded in 2002 to protect additional giant panda habitat. GNR (33°35'-33°45'N, 107° 51' – 108° 01' E) covers an area of 135 km², and is located on the south slope of the Qinling Mountainsdirectly adjacent to the eastern boundary of FNR. The average annual temperature at GNR is 11.5 °C with elevation ranging between 1,150 to 2,574 m. Arrow bamboo (located > 1,800 m) and dragon-head bamboo (located below 1,800 m and close to water) are common in the reserve. Because the GNR area was previously part of a state-owned forest enterprise, logging activities were a common source of disturbance. After the establishment of the Natural Forest Conservation Program in 1998, logging activities were banned (Liu et al., 2008). Giant pandas were recently spotted several times in dragon-head bamboo forest during spring and autumn in this reserve (Liu et al., 2013).

Sample processing

We collected bamboo samples from 57 plots across the FNR and GNR in April, July, and October 2013 (fig. 1). We chose these sampling periods because they are in the middle of spring (March to May), summer (June to August), and autumn (September to November). We divided the elevation into three strata: low elevation

stratum (1,500-1,750 m) with wooden and dragon-head bamboo, transition elevation stratum (1,750-2,200 m) with arrow and wooden bamboo, and high elevation stratum (2,200-2,700 m) with only arrow bamboo. We took wooden bamboo, arrow bamboo and dragon-head bamboo samples in Sanguanmiao, Guangtoushan, and Xigou where there were large forests of the three bamboo species. Each season we sampled one transect (arrow bamboo) at high elevation, two transects (arrow and wooden bamboo) at the transitional elevation, and two transects (wooden and dragon-head bamboo) at the low elevation. Each transect contained three or four plots. In each plot (1 x 1 m) we took samples (200–300 g, wet) of shoots, leaves, branches, one-year culms, and culms older than one year. Each sample was washed with distilled water, ground to 80 g mesh by a high-speed grinder, and dried to a constant weight at 60°C. We analyzed seven mineral elements (Cu, Zn, Fe, Mn, K, Ca, and Mg) by the standard atomic absorption spectrophotometry procedure (Skoog and West, 1980).

Data analysis

We tested the correlation between the seven elements using Pearson correlation and the mean content difference between levels of each factor (seasons, elevations, parts, and species) by non–parametric Kruskal–Wallis test (significance level at α = 0.05).

During spring and winter, giant pandas are present only at low elevations and absent at transitional and high elevations. In late spring, giant pandas leave low elevation areas and migrate rapidly through the transitional region to reach high elevation areas where they stay until the end of summer (Liu, 2001; Nie et al., 2014). To examine the relationships between mineral elements and spatial distribution of giant pandas, we summarized 35 variables based on the mean nutrient concentration in all bamboo parts (5 parts x 7 elements).

We used a Random Forest (RF) approach to model the presence and absence of giant pandas using these 35 predictive variables. We implemented the Random Forest package (Liaw and Wiener, 2002) in R software (RDevelopment, 2008) to build the models. RF is a non-parametric ensemble tree model for classification that operates by constructing a multitude of decision trees. RF can accommodate large number of input variables without overfitting and it does not require a specific distribution for predictive variables (Breiman, 2001; Svetnik et al., 2003; Prasad et al., 2006). The RF model is particularly well suited for modeling environmental co-variables which are often correlated (Breiman, 2001; Svetnik et al., 2003; Prasad et al., 2006). The algorithm of RF starts with the selection of bootstrapped samples from the original data. There are approximately 63% of the data in each bootstrap sample. Each decision tree grows with one bootstrap sample and a randomly selected subset of variables. Observations outside the bootstrap sample, called 'out-of-bag' observations, are used as testing data to examine the prediction error. One of the advantages of the RF model is that it provides variable importance by measuring how much model performance (specifically measured by the Gini index) declines if the variable is randomly permuted. The greater the increase

of prediction error, the greater the variable importance in the model. The final prediction is obtained by aggregating over the ensemble trees (Breiman, 2001; Cutler et al., 2007; Biau, 2012).

For a small number of plots, we did not collect oneyear culms or shoots because the two parts were not observed year-round. In order for RF to incorporate variables with missing data, we used multiple imputation by chained equation (MICE) (Buuren and Groothuis-Oudshoorn, 2011) to create plausible values for the missing data (Van Buuren and Oudshoorn, 1999). Compared with other methods, such as single imputation, MICE allows for the imputation of multivariate data based on the distribution of observed data, without the need to specify a joint distribution of predictor variables (White et al., 2011). It can also handle different variable types since each variable is imputed using its own imputation model, with good quality prediction and less biased estimates (Ambler et al., 2007). Normally, for a given missing data point, a multiple imputation method generates 3-5 imputations and missing values are replaced by the average of the multiple imputed values (Hui et al., 2004). In our research, we generated the imputed values using the MICE package in R software.

To take into account the varying availability of bamboo shoots over different seasons, we used the realized nutrient concentration of each plant part, that is, the product of measured or imputed nutrient values and availability of the specific part over the season for the predictive variables in the RF model. For species in which bamboo shoots are only available for a short period (spring for wooden bamboo and summer for arrow and dragon-head bamboo), we set the availability as 1 for the seasons when shoots were present and 0 for the other seasons. For the other species, the availability of leaves, branches, culm and one-year culm were set to 1 for all seasons. We next compared the mean difference of each mineral value between giant panda presence and absence for each predictor using the Wilcoxon rank sum test (significance level at α = 0.05). Finally, we built RF models for spring (SPR), summer (SUM), and autumn (AUT) to classify the presence and absence of giant pandas based on the nutrient concentrations determined for each element. To determine how the predictors associated with a suitable habitat for giant pandas in all three seasons, we built a combined model (COM), including all mineral data.

Results

Correlation of seven mineral elements

Most elements had a correlation value under 0.4 (table 1). Ca and Mg showed the strongest correlation (r = 0.69), followed by Cu:Zn (r = 0.64), Mn:Mg (r = 0.46), and Mn:Ca (r = 0.43) (fig. 1s) and lowest correlation was between Cu and Mg (r = 0.02).

Mean mineral difference test

We tested the mean difference between different levels for each factor (season, elevation, plant part, and

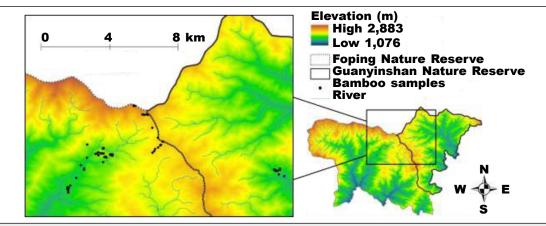


Fig. 1. Schematic map of the study area and sampling points.

Fig. 1. Mapa esquemático de la zona de estudio y los puntos de muestreo.

species) (see supplementary material). No significant differences were found between three elevation ranges for any mineral element (all p > 0.05). However, concentrations between the three seasons differed (all $p \le 0.05$) for all mineral elements except K. The highest concentrations of Cu and Zn were in spring, followed by those in autumn and summer (fig. 2, table 1s), while Fe had a higher value in summer than spring and autumn (table 1s). For Mn, Ca, and Mg, samples collected in autumn were at a highest concentration followed by those in spring and summer.

Within species, only Mn significantly differed between the three bamboo species (p = 0.028), with wooden bamboo showing the highest level, followed by dragon—head and arrow bamboo (fig. 2). Finally, all mineral elements demonstrated a significant difference between the five plant parts (all $p \le 0.05$). The highest concentration of K was found in shoots; Mn, Ca and Mg were in highest concentration in leaves; Cu, Zn and Fe were at their highest levels in branches (table 1s).

In SPR, Zn found in branches, Ca found in shoots, and Mn found in one—year culms and in branches were all in higher concentration in habitats where giant pandas were present than in habitats where they were absent (fig. 3). In SUM, the Fe, Ca and Mg content of shoots was higher in habitats where giant pandas were present. However, Cu, K, Ca, and Mg concentrations found in leaves were significantly higher in habitats where giant pandas were absent (fig. 3). In AUT, Mn concentration found in leaves and one—year culms was higher in habitats where giant pandas were present, while Mg concentration found in branches was higher in habitats where giant pandas were absent (fig. 3) (all $p \le 0.05$).

Classification models

Among the four models, the lowest error rate occurred in SUM (10.5%), while the highest error rate was 31.6% in AUT. COM and SPR had 20.4% and 18.6% error rates, respectively. In COM, Mn found in

one—year culms had the largest decrease of the Gini index, followed by Mn in culms, Ca in shoots, and Mn in leaves (fig. 4). In spring, Mn found in one—year culms showed the highest variable importance, followed by Ca in shoots, and Ca and Mn in branches. However, Mg and Ca found in shoots and K and Mg found in leaves were the most important predictors in summer. Mn and K found in one—year culms, Mg found in branches, and Mn found in culms had the highest importance measure in AUT (fig. 4).

Discussion

Our results show strong associations between giant pandas' habitat selection and bamboo mineral content. The results show contrasting dietary consequences as giant pandas migrate to different habitats at different times of the year (fig. 3). Our results suggest that the nutrient composition in the giant pandas' forage

Table 1. Correlation of seven bamboo mineral elements.

Tabla 1. Correlación entre siete elementos minerales del bambú.

	Cu	Zn	Fe	Mn	K	Ca
Zn	0.64					
Fe	0.36	0.19				
Mn	0.26	0.34	0.11			
K	0.19	0.07	0.13	0.11		
Ca	0.36	0.36	0.06	0.43	0.29	

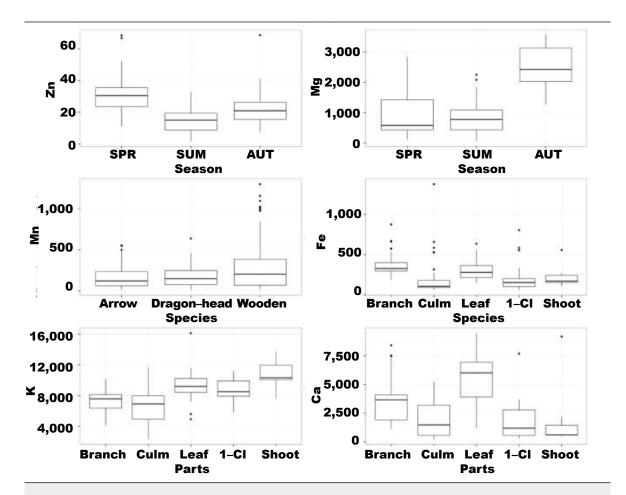


Fig. 2. The effect of different factors on mineral content (mg/kg). The box plots show the distribution of six mineral content values differentiated by levels of three factors (seasons, bamboo species, and bamboo parts). We only show here the elements with the most significant differences between levels: 1–Cl, one year culm.

Fig. 2. Efecto de diferentes factores en el contenido de minerales (mg/kg). Los diagramas de caja muestran la distribución de los valores del contenido de seis minerales según la categoría de tres factores (estación, especie de bambú y partes del bambú). Solo mostramos aquí los elementos con las diferencias más significativas entre categorías: 1–Cl, cañas de un año.

bamboo is an important co-variant as they migrate seasonally, and such nutrient differences can be used to differentiate giant pandas' seasonal habitat selection. Our study supports the hypothesis that the seasonal migration pattern might have evolved to facilitate access to specific composition of nutrients from forage.

One of the important hypotheses previously proposed to explain giant pandas' movement is that shoots sprouting in the Qinling Mountains in late spring and early summer drives their seasonal movements from low to high elevation (Pan et al., 1988). Evidently, dragon—head and arrow bamboos produce shoots only during summer and wooden bamboo shoots are available only in spring (Nie et al., 2014). Additionally, for almost all bamboo species, due to temperature

differences, the shoots sprout first at lower elevation and shift to higher elevation sequentially. This coincides closely with giant pandas migrations to higher elevations. However, this hypothesis is mostly based on observed shoot availability and it is unclear how mineral content in bamboo shoots associates with such patterns.

By incorporating the availability of shoots with the measured mineral content, our study allowed us to quantitatively test for associations between panda habitat choice and various minerals in shoots. Based on our classification model, we found that the concentrations of Fe, Ca, and Mg in shoots effectively predict differences in presence and absence at habitats during the summer. The highest concentrations of K, Ca, and Mg in summer were found in bamboo leaves

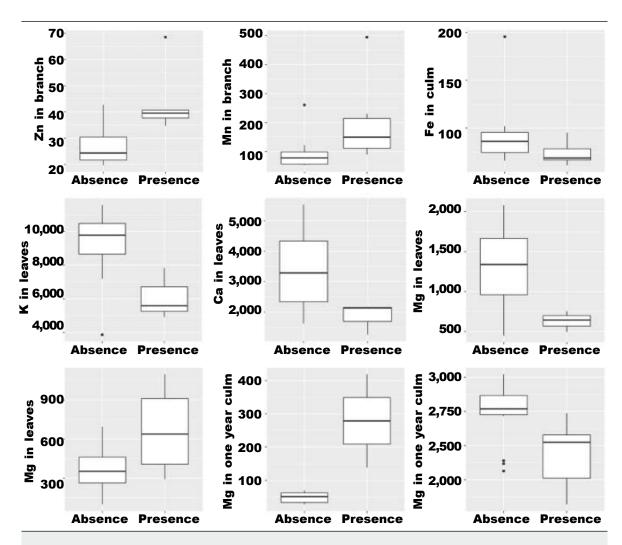


Fig. 3. The contrast of nutrient concentration (mg/kg) between the seasonal habitat where giant pandas are present and the habitat where they are absent. The concentration of nine nutrients in spring (top row), summer (middle row), and autumn (bottom row) and from different bamboo parts are compared.

Fig. 3. Diferencia de concentración de nutrientes (mg/kg) entre el hábitat estacional en el que el panda gigante está presente y el hábitat en el que está ausente. Se compara la concentración de nueve nutrientes en primavera (fila superior), verano (fila media) y otoño (fila inferior) y en diferentes partes del bambú.

in areas where giant pandas were absent (fig. 3). It is possible that shoots are preferred because they have the lowest concentration of fiber and cost less energy to obtain than other parts of the plant.

In our results, Mn was the most influential predictor of giant pandas' presence not only in COM but also in SPR and AUT. Lack of Mn can lead to many physiological problems in animals, e.g., decrease of sperm, fecundity decline, premature birth and lactation cessation (Kemmerer et al., 1931; Plumlee et al., 1956; Hurley and Doane, 1989). The physiological function of Mn could contribute to its importance, especially when giant pandas are in estrus in spring and pregnant in summer (Schaller et al., 1989; Zhu et al., 2001). Similarly, Ca is mainly stored in bones, and

lack of Ca could cause arrested development, bone deformities, and miscarriages or stillbirths (Hightshoe et al., 1991; Bhanderi et al., 2014). Animals will also experience face muscle twitching or convulsions if they cannot get enough Mg (Kaneko et al., 2008; Chandra et al., 2013). The physiological function of Ca and Mg might have contributed to the observations that Ca and Mg were more concentrated in the presence areas in summer and they were ranked high in our models. Ultimately, if giant panda migration is evolutionarily driven by nutrient gradients, our models suggest that Ca, Mg, and Mn play an important role in the process. Further research is required to investigate the specific biochemical interactions of the mineral elements in panda physiology.

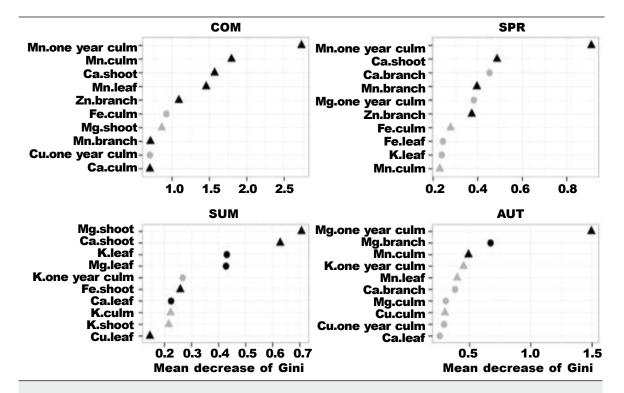


Fig. 4. Relative importance of variables in four Random Forest models: COM, combined model; SPR, spring model; SUM, summer model; and AUT, autumn model. Variable importance plots show the mean decrease of Gini index of the top 10 influential predictors in four RF models: the black symbols indicate a significant difference between habitats where giant pandas were present and absent and the grey symbols indicate no significant difference; triangles indicate the mean of the presence habitat is higher than that of the absence habitat and the opposite is represented by circles.

Fig. 4. Importancia relativa de las variables en cuatro modelos de bosques aleatorios: COM, modelo combinado; SPR, modelo primavera, SUM, modelo verano; AUT, modelo otoño. En los gráficos de la importancia variable se muestra la reducción media del índice de Gini de los 10 factores de predicción más influyentes en cuatro modelos de bosques aleatorios: los símbolos negros indican que hay una diferencia significativa entre los hábitats donde el panda gigante estaba presente y ausente, y los símbolos grises indican que no hay diferencia significativa entre ambos; los triángulos indican que el promedio del hábitat donde estaba presente es superior al del hábitat donde no lo estaba; lo contrario se representa con círculos.

Our comparison of bamboo samples between presence and absence habitats provides a new approach to identifying potentially important dietary factors underlying panda movements. Using this mean comparison method, we found that predictors such as K and Mg found in leaves in SUM and Mg found in branches in AUT with high variable importance were significantly lower in the elevations where giant pandas were present. We speculate that despite the low concentration of these mineral elements in the presence habitat of giant pandas, the level may still meet their requirements for that mineral, and giant pandas may prefer to acquire them from other parts of bamboo. On the other hand, other predictors such as Ca found in branches in SPR, K found in one-year culms in SUM, and K found in one-year culms in AUT did not show significant differences between presence and absence habitats, but they ranked high in their importance for classifying the presence of giant pandas. This result suggests small but consistent differences in content between presence and absence habitats. It could also indicate that non–linear relationships exist between these habitats (fig 4). Further studies are needed to examine those relationships and how they affect giant panda activities (Swaisgood et al., 2009).

The RF model can perform well using predictors with correlations and still provide an accurate variable importance ranking (Breiman, 2001; Biau, 2012; Freeman et al., 2012). Normally in linear model, the proper way to deal with correlated variables having redundant information is to remove one or multiple correlated variables. In our case, all elements are included because they have specific physiological functions. For example, Ca and Mg had the highest correlation and they demonstrated similar importance rankings in each model. Our model highlights the advantage

of a non-parametric RF model which often deals with a large number of correlated nutrient variables.

It is worth noting that this study examined the total concentration of key nutrients. It does not consider the fiber-bonded mineral element. As giant pandas' digestive system has lower efficiency absorbing fiber-bonded minerals than non-fiber-bonded minerals (Dierenfeld et al., 1982), the portion of fiberbonded mineral element in bamboo is an important factor that affect the amount of nutrient utilized by giant panda (according to analysis done on leaves of Phyllostachys aureosulcata, 25 % of the total mineral are fiber bound and not available to giant pandas (unpublished data and personal communication with Dr. Michael Power). However, measurements of total mineral content are a necessary first step to understanding the patterns of giant panda habitat selection with the nutrient landscape. Knowing what minerals are concentrated where within the landscape and how the concentration of these elements changes both spatially and temporally improves our understanding of potential drivers of giant panda migration and habitat preferences. This knowledge also provides insights for improving conservation planning and management.

With the increasing giant panda density in FNR there has been an increasing number of observations of giant pandas dispersing from west to east across the reserve boundary into GNR to colonize unoccupied habitat (Hu et al., 2010; Liu et al., 2013). In order to create a suitable environment, local governmental and conservation organizations have been planting bamboo in GNR (World Wide Fund for Nature, 2006). Our results will help to provide guidelines for determining the bamboo species and the locations for the planting practices. According to our results, wooden bamboo is recommended for giant panda habitat restoration in low elevations because of its highest concentration of nutrients. GNR does not have any large distribution of wooden bamboo due to its logging history. We recommend planting wooden bamboo in GNR, especially in low elevational areas, because it should support giant panda dispersal from FNR in the west to GNR in the east. To achieve better results, bamboo should be planted every few years to minimize the impact of periodic flowering on the panda population (Kawamura, 1927; Chai et al., 2006). In 2011 and 2013 our team started planting wooden bamboo in GNR. However, bamboo contains the lowest nutrient levels in summer, and only arrow bamboo is distributed in high elevation. Ensuring adequate arrow bamboo in high elevational areas is critical to giant panda conservation because an abundant food supply can compensate for lower nutrient content. Since the establishment of the reserves, economic activities have been banned within them. However. some wild giant pandas still live in the area outside the protected areas where they have to compete with local people harvesting bamboo shoots. We strongly recommend that future conservation policies are enacted to develop a sustainable bamboo shoot harvesting plan to ensure a sufficient supply of shoots for giant pandas both inside and outside reserves.

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Supplementary material

Table 1s. Mean concentration (mg/kg) and p-value of each factor for seven mineral elements: * p-value < 0.05. (Season: SPRm, spring mean; SUMm, summer mean; AUTm, autumn mean. Elevations: H, high; L, low; T, transition. Species: Wb, wooden bamboo; Dhb, dragon head bamboo; Ab, arrow bamboo. Part: Sh, shoots; Lv, leaves; Br, branches; Cl, culms; 1-Cl, one-year culms).

Tabla 1s. Concentración media (mg/kg) y valor de p de cada factor para siete elementos minerales. * p < 0,05. (Para las abreviaturas, véase arriba).

Season								
	<i>p</i> –value	SPRm	SUMm	AUTm				
Cu	*	17.44	9.05	10.04				
Zn	*	30.95	14.51	22.16				
Fe	*	232.30	353.36	208.90				
Mn	*	230.64	132.11	320.00				
K		8,158.59	7,620.37	8,116.10				
Са	*	3,858.59	1,824.48	4,345.19				
Mg	*	863.79	823.77	2,524.31				

Elevation			
<i>p</i> –value	Н	L	Т
	12.66	12.11	10.94
	21.61	22.84	20.67
	320.76	253.46	263.04
	184.37	291.26	178.06
	8,305.18	7,878.96	7,826.25
	3,044.40	3,092.10	3,492.78
	1,322.32	1,313.33	1,549.34

Species								
	<i>p</i> –value	Wb	Dhb	Ab				
Cu		12.01	10.29	12.05				
Zn		23.67	18.95	20.79				
Fe		239.87	255.50	309.34				
Mn	*	295.29	176.48	167.46				
K		7,968.46	7,793.12	7,971.04				
Са		3,537.71	2,579.12	3,232.01				
Mg		1,567.96	1,140.42	1,366.30				

Part					
p-value	e Sh	Lv	Br	CI	1–Cl
*	14.47	14.04	15.55	7.02	8.55
*	20.11	23.84	27.05	17.70	15.43
*	218.71	302.38	369.33	183.22	211.64
*	61.29	381.24	208.93	178.06	78.16
*	10,740.36	9,258.57	7,225.72	6,618.28	8,743.27
*	1,869.84	5,430.65	3,536.32	1,797.73	1,826.67
*	925.52	2,113.83	1,406.93	1,078.93	871.35

Table 2s. Mean concentration (mg/kg) and p-value of each RF model for every predictor: * p-value < 0.05; - not applicable: Prm, presence mean, Abm, absence mean: 1–Cl, one-year culms.

Tabla 2s. Concentración media (mg/kg) y valor de p de cada modelo de bosque aleatorio para cada factor de predicción: * p < 0,05; – no aplicable. (Para las abreviaturas, véase arriba).

_		COM			SPR			SUM			AUT	
	val	ue Prm	Abm	p–va	lue Prm	Abm	<i>p</i> –valı	ue Prm	Abm	<i>p</i> –va	ue Prm	Abm
Cu.shoots		5.93	3.52		11.13	4.40		7.62	5.66	_	_	-
Zn.shoots		6.91	3.96		12.75	4.81		9.41	6.44	_	_	-
Fe.shoots		106.74	52.70		117.86	54.83	*	329.84	91.03	_	-	_
Mn.shoots		22.17	17.76		36.20	16.61		41.15	31.72	_	-	-
K.shoots		4,937.97	3,073.36	3	7,431.86	3,446.5	4	10,640.83	5,168.47	7 <u> </u>	-	-
Ca.shoots	*	1,437.86	251.45	*	2,812.09	400.03	*	1,586.33	356.46	_	-	-
Mg.shoots		600.14	174.94		998.56	319.45	*	1,070.83	224.86	_	_	_
Cu.leaves		16.92	12.94		25.68	26.27	*	14.96	8.04		9.00	9.48
Zn.leaves		25.78	23.02		32.28	34.95		18.06	15.96		22.60	23.47
Fe.leaves		305.75	300.25		324.79	410.69		433.23	295.27		232.08	224.05
Mn.leaves	*	592.32	283.11		674.53	392.80		227.92	159.75		666.29	365.32
K.leaves	*	8,476.56	9611.15		9,603.23	10779.7	1 *	6,117.00	9,833.75	5	8,361.14	8,437.92
Ca.leaves		5,636.73	5,350.61		5,983.42	6311.95	· *	1,852.00	3,494.22	2	6,912.07	7,104.79
Mg.leaves		2,142.04	2,099.35	5	1,602.67	1,752.38	3 *	630.33	1,362.13	3	3,329.29	3,342.54
Cu.branche	s	17.60	14.76		23.39	24.56		9.32	9.35		15.36	14.61
Zn.branches	s *	34.48	23.79	*	41.89	29.79		15.70	16.86		35.13	28.53
Fe.branches	S	330.54	390.18		297.52	369.50		394.43	448.89		336.19	327.41
Mn.branche	s *	274.64	182.96	*	226.52	102.81		190.71	169.64		358.73	260.84
K.branches		7,546.93	7,065.38	3	7,022.11	6,538.24	1	7,473.33	6,542.34	ļ	8,103.29	8,158.13
Ca.branche	s	3,764.07	3,439.22	2	3,765.60	3,994.5	1	1,798.17	1,718.8		4,605.07	5,316.63
Mg.branche	S	1,392.61	1,401.47	7	562.77	679.73		909.17	827.88	*	2,429.64	2,707.58
Cu.culms		7.38	7.01		6.50	5.17		10.11	8.56		7.10	6.31
Zn.culms		19.90	16.93		24.57	27.08		16.99	12.74		16.48	14.91
Fe.culms		180.19	203.41		125.11	86.27		454.73	335.24		117.61	115.50
Mn.culms	*	313.86	113.18		229.96	92.54		167.20	98.58	*	460.62	148.13
K.culms		6,945.27	6,322.02	2	5,876.52	6,607.14	1	7,396.67	5,083.78	3	7,820.57	7,759.17
Ca.culms	*	2,341.86	1,506.31		2,603.95	1,512.9	5	2,512.33	776.66		2,006.71	2,474.2
Mg.culms		1,117.65	966.87		369.86	419.84		1,072.33	469.06		1,884.86	2,040.88
Cu.1–Cl		7.99	8.78		7.60	8.13		6.98	8.29		8.81	9.91
Zn.1–Cl		16.64	15.73		17.25	21.54		13.93	11.95		17.18	16.42
Fe.1–Cl		217.71	192.58		144.66	93.12		437.88	277.61		196.40	153.80
Mn.1–Cl	*	140.56	57.56	*	74.57	29.79		148.38	66.70	*	203.19	66.21
K.1–Cl		8,590.33	8,883.44	ļ	8,885.35	9,611.32	2	7,991.67	9,078.19)	8,551.87	8,077.8
Ca.1–Cl		2,340.55	1,755.04	ļ	2,839.74	2,407.89)	3,261.33	1,295.57	7	1,446.73	1,878.0
Mg.1–Cl		1,008.96	896.07		537.42	659.70		996.33	527.02		1,485.91	1 565 4

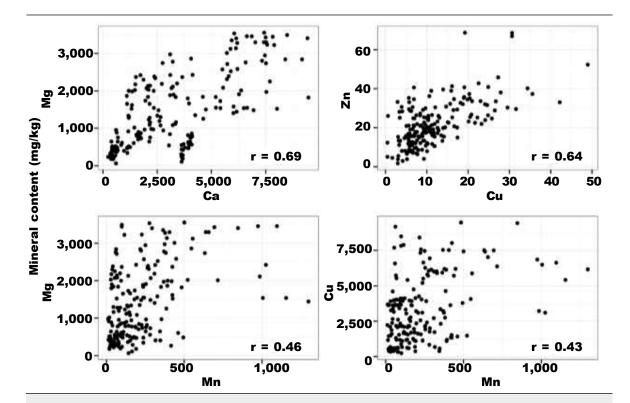


Fig. 1s. Correlation of mineral elements. Scatter plots show the four pairs of the mineral elements with the strongest correlation.

Fig. 1s. Correlación de elementos minerales. En los gráficos de dispersión se muestran los cuatro pares de elementos minerales con la mayor correlación.