

ISOTOPIC CHARACTERIZATION OF *CEDRELA* TO VERIFY SPECIES AND REGIONAL PROVENANCE OF BOLIVIAN TIMBER

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ABSTRACT

With increasing concerns about sustainable exploitation of tropical timber, there is a need for developing independent tools to check their origin. We evaluated the potential of tree-ring stable isotopes for identifying four *Cedrela* species (*C. balansae*, *C. fissilis*, *C. odorata*, and *C. saltensis*) and for identifying geographic origin of *C. fissilis* and *C. odorata*, two of the most intensively exploited species. We studied differences in $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ of wood among 11 forest sites (163 trees). We quantified isotope composition of 10-year bulk samples, and for a subset we also evaluated isotopic annual fluctuations for the last 10 years. Although annual isotopic variability was not correlated to precipitation or elevation, we found a significant relationship between the 10-year bulk stable-isotope composition and average precipitation and elevation. However these relationships were not consistent across all sites. We also explored isotopic site and species differentiation using Kernel Discriminant Analyses. Site discrimination was low: 30% accuracy for *C. odorata*, and 40% for *C. fissilis* sites. However, species discrimination was 57.5% for *C. odorata* and 95.3% for *C. fissilis*. These results suggest that although $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ isotopic analyses hold potential to verify species identification, discrimination of geographical origin within a country may still be challenging.

Keywords: Illegal logging, *Cedrela*, stable isotopes, discriminant analysis, tropical timber.

INTRODUCTION

Illegal logging practices lead to loss of natural resources, community conflicts and economic problems. Illegal low-priced timber not only damages local and global forest markets (Abt Associates Inc. 2006; Blaser 2010), but also threatens sustainable management of tree species and forest areas (ABT 2017). The value of the wood trade associated with illegal logging in South American countries (Brazil, Colombia and Peru) has reached \$682 million USD, with illegal logging rates ranging from 53 to 75% (Jianbang *et al.* 2016). The most common type of illegal timber trade is the fraud concerning

false declarations of sites of origin based on falsified documents (Lowe *et al.* 2016). Increasing sustainability concerns call for developing tools to independently verify that the origins of timbers are in compliance with national and international regulations (Degen and Fladung 2007). A technique based on the intrinsic characteristics of the wood could help confirm the origin of the timber more reliably and thus help combat illegal timber trade.

The cellulose produced by trees to form wood tissue records the carbon and oxygen isotopic composition of the CO_2 and H_2O taken up by trees during photosynthesis and water uptake. These isotopic signatures can reveal important information about the environmental conditions the tree was experiencing at the time of growth. During photo-

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synthesis, plants discriminate against ^{13}C . The discrimination has been modelled by Farquhar *et al.* (1982) and is a function of the ratio of intercellular to ambient CO_2 concentration. This strongly correlates with water-use efficiency (Farquhar and Richards 1984) that can be estimated using tree-ring archives (van der Sleen *et al.* 2015). The oxygen isotopic composition ($\delta^{18}\text{O}$) in cellulose of tree rings records the isotopic composition of rainfall (Brienen *et al.* 2012). Recent studies have reported the relation between stable isotopes in Amazonian trees and precipitation (Evans and Schrag 2004; Jenkins 2009; Brienen *et al.* 2012; van der Sleen 2014). Stable isotopic composition of rainfall not only correlates to the amount of rain, it also shows distinct geographical patterns (Rozanski *et al.* 1993; West *et al.* 2010). Furthermore, both $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ are also correlated with elevation (Körner *et al.* 1988; Rozanski *et al.* 1993; Körner 1998), which could result in distinct isotope landscapes that can be useful to trace timber origin.

Stable isotopes have become an important tool to trace geographical origin of many biological materials. In particular, its application to ecology enables researchers to trace origin and migration of animals (Hobson 1999; Hobson and Wassenaar 1999; Rubenstein and Hobson 2004). Stable isotope analyses have also been successfully used in forensic applications. For example, stable isotopic composition of human hair is linked with geographic origin as it shows strong correlations with the geographic distribution of drinking water isotopic composition (Ehleringer *et al.* 2008).

In recent years, the analysis of stable isotopes has proven to be useful to accurately determine the origin of foods (Versini *et al.* 1997; Rossmann 2001; Oulhote *et al.* 2011; Driscoll *et al.* 2020) and drugs (Ehleringer *et al.* 2000; Booth *et al.* 2010). Geographic origin tracing by stable isotopes has also been tested for timber by Boner *et al.* (2007), Förstel *et al.* (2011), Förstel and Hützen (1983), Kagawa and Leavitt (2010), and Vlam *et al.* (2018). Although some of these studies have shown the potential for geographic identification at small scale (>14km) (Vlam *et al.* 2018; Watkinson *et al.* 2020), these studies reached different conclusions on the potential of timber tracing across larger spatial scales.

Stable isotopes have also shown promise for species identification (Chouvelon *et al.* 2014; Hamer *et al.* 2015), being successfully applied to fishes (Oliveira *et al.* 2011), birds (Militão *et al.* 2014), frogs (Dittrich *et al.* 2017) and archaeozoological material (Gorlova *et al.* 2015). To date, stable isotopes have not yet been used to identify tree species in a forensic context. Our study represents a first exploration on the potential of stable isotopes for the identification of *Cedrela* species and tracing back their site of origin.

Cedrela is an intensively harvested and traded tropical timber genus (Richter and Dallwitz 2000), widely distributed across climatic and environmental gradients throughout tropical America and the Caribbean islands (Wagenführ 2007), from moist to dry tropical forests and along a wide altitudinal range (Mostacedo *et al.* 2003; Navarro 2011; Navarro-Cerrillo *et al.* 2013). *Cedrela* has been traded as roundwood, lumber, veneer and exported predominantly to Europe and Northern America (Wagenführ 2007). Overexploitation of *Cedrela* species has caused population declines in recent decades (Mostacedo and Fredericksen 1999, 2001), motivating the Convention on International Trade in Endangered Species of Wild Fauna and Flora (CITES) to include *Cedrela* species in its Appendix II (CITES 2019). Providing an objective, fast to implement, and reliable way to differentiate between *Cedrela* species and sites of origin across Bolivia, such as using isotopic compositions based on $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$, would greatly help in regulating and preserving these valuable and vulnerable tree species.

We expect isotopic signatures of carbon and oxygen to vary between sites, particularly at higher elevations, where spatial differences in terms of precipitation and topography are larger compared to the lowlands (Körner *et al.* 1988; Rozanski *et al.* 1993; Körner 1998; Brienen *et al.* 2012). Our research questions are: (1) Can we identify the geographic origin of *Cedrela odorata* and *C. fissilis* based on their isotopic signatures? (2) Can the relationship between precipitation, elevation and isotopic ratios be used to predict the likelihood of timber coming from specific sites? (3) Are the isotopic signatures different among *Cedrela* species? And, if so, to what extent do they differ? To address these questions, we collected *Cedrela* samples across multiple Bolivian forest types and assessed the isotopic

variation within species in relation to precipitation, elevation and site of origin.

METHODS

Sampling Sites

We performed species identification analysis on four *Cedrela* species (*C. balansae*, *C. fissilis*, *C. odorata*, and *C. saltensis*), and more-detailed origin identification on two of them (*C. fissilis* and *C. odorata*). Wood samples from 163 trees (2 wood cores per tree) were collected from eleven natural forest sites distributed across the species distribution areas in Bolivia (Figure 1). Cores were collected at breast height (130 cm) using a 5-mm-diameter increment borer (Haglöf Inc., Sweden). We targeted large/dominant trees in each site, as there is a height gradient associated with leaf $\delta^{13}\text{C}$ values caused by the access to light and humidity (Ometto *et al.* 2002). Average tree diameter was 41 cm for both *C. fissilis* (range: 18–103 cm) and *C. odorata* (13–103 cm). Within each site, sampled trees were separated by at least 3 m for *C. fissilis* and 85 m for *C. odorata*. We tried to cover most of the species distributions in Bolivia and thus minimum distances between different sampling sites were 283 to 421 km for *C. odorata*, and 78 to 400 km for *C. fissilis*. We also sampled trees between sites, along roads and in villages, to improve spatial coverage and capture gradual spatial variation between sampling sites.

When species identification was not possible directly in the field, we collected botanical samples that were later identified by Alejandro Araujo Murakami, at the Museo de Historia Natural Noel Kempff Mercado (Bolivia). This was done for 27% of the sampled trees. Climate data were obtained from the reporting agency of Servicio Nacional de Meteorología e Hidrología (SENAMHI 2018; Table 1).

Samples were air dried and carefully polished beginning with ISO P40-grit (425–500 μm) sandpaper and using progressively finer sizes until ultimately sanding with ISO P600-grit (24.8–26.8 μm) (Orvis and Grissino-Mayer 2002). *Cedrela* species have clearly visible, ring-porous tree rings, with parenchyma bands as ring boundaries (Worbes 1999; Paredes-Villanueva *et al.* 2016). Tree-ring boundaries were identified using a binocular mi-

croscope (Leica S6E) coupled to an LED light source. Annual ring formation in *Cedrela* species is well established (Worbes 1999; Dünisch *et al.* 2002; Brienen and Zuidema 2005; Bräuning *et al.* 2009; Paredes-Villanueva *et al.* 2016). Tree rings were assigned to the calendar year in which their growth started, because growth season in these forests typically lasts from September of the current year to August of the following year (Schulman 1956). Next, annual ring widths within each increment core were measured, compared, and crossdated using TSAP/LINTAB software/hardware combination (Frank Rinn, Heidelberg, Germany) to a resolution of 0.01 mm. Missing and false rings, suggested by crossdated samples, were visually identified, re-measured and checked in an iterative process. We also estimated the mean first-order autocorrelation of raw tree-ring width data, a measure of the year-to-year growth similarity (AR1); mean intercorrelation among series, quantifying the similarity in width among trees (r); and mean sensitivity, which measures the year-to-year variability in width of consecutive rings (MS) for each site (Supplementary Material Table S1) using the dplR 1.7.2 (Bunn *et al.* 2021) R package. However, because the analyzed time-series spans were short (10 years), the results should be interpreted with caution.

Isotope Analyses

We conducted exploratory analyses of $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ stable isotopes series of 10 years in bulk and annual samples separately. Wood flakes were sliced from the outermost 10 rings pooled into one sample (10-year bulk samples) of each core using a scalpel and microtome. Wood material from separate annual tree rings was also sampled from 3–4 trees per site (10% of the sample set) for all the sites, except Yapacaní. Crude cellulose was extracted following the adaptation of the Jayme-Wise method (Wieloch *et al.* 2011) described in Vlam *et al.* (2018) and then oven-dried at 50°C. For $\delta^{13}\text{C}$ analysis, cellulose samples were combusted on an element analyzer, and for $\delta^{18}\text{O}$, cellulose was pyrolyzed using a glassy carbon reactor at 1450°C. Both were coupled to a continuous flow isotope ratio mass spectrometer (Sercon Hydra 20–20). The values of the isotopes ratios ($\delta^{13}\text{C}$ and $\delta^{18}\text{O}$) per site were measured in parts per thousand (‰) according to the

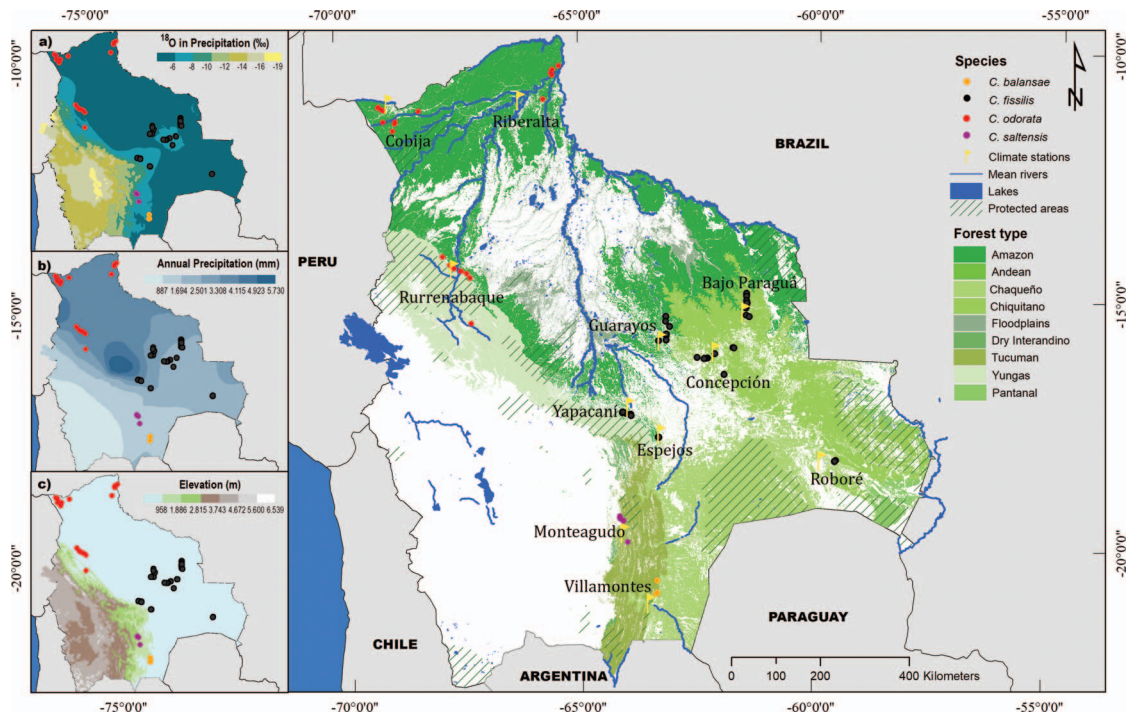


Figure 1. Distribution of *Cedrela* sampling sites and relevant environmental variables in Bolivia. Forest cover: Autoridad de Bosques y Tierra, 2015. (a) $\delta^{18}\text{O}$ spatial variation in mean annual precipitation (Bowen and Revenaugh 2003), (b) Mean annual precipitation of the last eleven calendar years (University of East Anglia Climatic Research Unit 2008), and (c) Digital elevation model (NASA/METI/AIST/Japan SpaceSystems and U.S./Japan ASTER Science Team 2019).

Table 1. *Cedrela* species and number of samples collected from 11 study sites in Bolivia.

Species	Sites	# Trees with 10-year Bulk Samples	# Trees with Annual Samples	# Trees with Vouchers	Average Elevation of Samples (m a.s.l.)	Annual Precipitation During Last 10 Years of Tree Growth (mm)	$\delta^{18}\text{O}$ in Precipitation (‰) ^c (vs SMOW)
<i>C. balansae</i>	Villamontes	10		3	588	^b	-6.4
<i>C. fissilis</i>	Bajo Paraguá	21	3	2	292	^b	-6.6
	Concepción	21	3	^a	468	1045	-7.3
	Espejos	21	3	^a	481	1552	-7.5
	Guarayos	21	4	13	224	1421	-6.6
	Roboré	21	3	13	635	1100	-6.3
	Yapacaní	10		5	318	1925	-7.5
<i>C. odorata</i>	Cobija	10		^a	274	2053	-5.8
	Riberalta	10		^a	145	1837	-5.9
	Rurrenabaque	10		4	309	2057	-6.8
<i>C. saltensis</i>	Monteagudo	8		4	1595	^b	-8.8
Total		163	16	44			

^aNo botanical samples were collected, but identification is based on previous collections.

^bNo climate data available from climate stations for the studied period.

^cInterpolated $\delta^{18}\text{O}$ variation in mean annual precipitation (Bowen and Revenaugh 2003).

international Vienna Standard Mean Ocean Water (VSMOW) for oxygen, and Pee Dee Belemnite (PDB) standards for carbon.

Statistical Analysis

Statistically significant differences between group means for species and sites were initially determined by one-way ANOVA. We then evaluated the relationship between isotopic composition, and local precipitation and elevation, using a mixed-effect modeling approach on annual measurements in *C. fissilis*, and on 10-year bulk measurements for *C. fissilis* and *C. odorata* with nlme 3.1–148 (Pinheiro *et al.* 2020) and MuMIn 1.43.17 (Bunn *et al.* 2021) R packages. We ran separate mixed models per species, to reflect potential species-specific responses. Sites were included in the models as random terms. In the case of annual time-series with multiple measurements per individual, tree identity was used as an additional random term. Annual precipitation of the last ten years was averaged for the studied sites. To be included in the 10-year bulk dataset, annual isotopic data were also averaged by the total years of measurement. We found no reliable climate data from Bajo Paraguá, and thus these samples were excluded from this analysis.

We performed a multi-step data analysis on the $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ data to evaluate whether this approach can consistently identify sites and species. As the dates of tree rings are generally not known in confiscated timber, we first needed to evaluate if taking into account annual variability may influence the main isotopic patterns and potentially hinder species or site identification. To test this, we compared two approaches for using the $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ annual data: (i) using raw averages of isotopic values, and (ii) using weighted averages based in their corresponding ring width. The idea behind the weighting correction is controlling for the fact that larger rings may contribute more to the mean isotopic value than small ones in the pooled sample. We performed a Pearson correlation to compare raw averaging and weighted averaging. After selecting the type of annual transformed data, these were then used as additional samples in the 10-year bulk dataset as input for the Kernel Discriminant Analysis (KDA). Pearson correlations between diameter at breast height (DBH) and stable isotope

values were also calculated to assess if there was an ontogenetic effect in the sampled trees. Statistical analyses were performed in R version 3.4.3 (R Development Core Team 2017), using the ggplot2 2.2.1 (Wickham 2016) and ggpubr 0.1.6 (Kassambara 2017) packages.

The site differentiation potential was assessed by a discriminant analysis on the $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ in *C. odorata* (10-year bulk dataset) and *C. fissilis* (10-year bulk plus average annual data in one set). The Kernel Discriminant Analyses were done with the package ks 1.10.6 (Duong 2007, 2017) in R version 3.3.3 (R Development Core Team 2017), and were based on randomized samples and variables in every run. The classification results allowed us to assess the site differentiation measured by the frequency in which each site is assigned to either a correct or erroneous class.

Assignment Tests

KDA separates samples based on an *a priori* classification assignment (to specific site classes) and looks for the optimal non-linear combination of variables (component loadings) for maximal separation of the samples in a two dimensional space (Baudat and Anouar 2000). KDA's learning algorithm uses Bayes discriminant rule (Duong 2007) and needs to be trained in order to assess its discrimination power. Therefore, our data were split in two sets: 80% for training and 20% for testing the model in each of the analyses. Smoothed Cross Validation (SCV) error (Duong 2007) was applied to test the correctness of the site assignments. KDA testing generated confusion matrices showing the frequency at which each site was wrongly classified. After 100 randomization runs, we checked with which site a single sample origin could be confused most. The final classification error was expressed in percentage (%), where a cross-validation error of 0% would indicate that all samples were correctly assigned to their respective origins. Finally, the average errors per site identification across the 100 runs were obtained together with their corresponding standard deviations. For the species discrimination, KDA tests were performed on the 10-year bulk data by using four species as classes (*C. balansae*, *C. fissilis*, *C. odorata* and *C. saltensis*) and following the same steps described above.

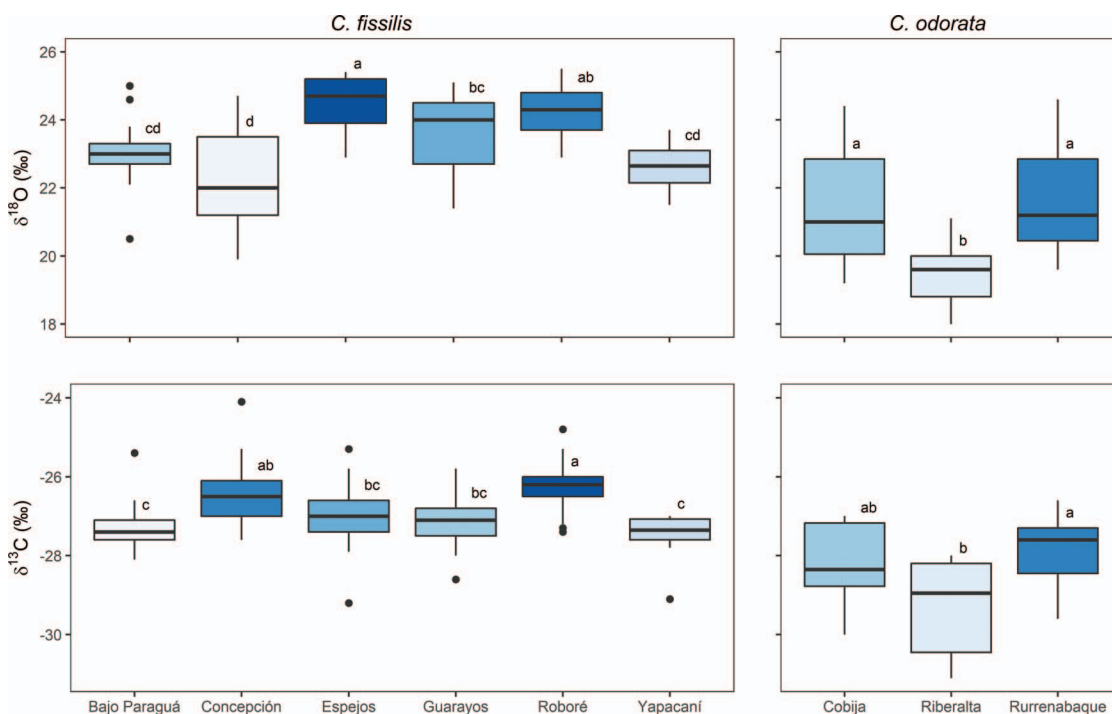


Figure 2. Variation of $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ isotopes composition in *C. fissilis* and *C. odorata* trees from 9 sites in Bolivia. Different lowercase letters above boxplots indicate statistically significant differences (one-way ANOVA, Tukey's HSD test, $\alpha = 0.05$) between the sites. Upper and lower box hinges indicate 25th and 75th quantiles, respectively, and horizontal lines within boxes indicate medians. Median differences are shown as a blue-based color gradient.

RESULTS

Raw $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ data suggest differences between sites for *C. fissilis* and *C. odorata* (Figure 2). Higher isotopic values were observed in *C. fissilis* trees from Espejos for $\delta^{18}\text{O}$ and Roboré for $\delta^{13}\text{C}$. *C. odorata* trees from Riberalta had the most contrasting isotopic signatures. Differences between other sites were smaller. Consistently, there were statistically significant differences between group means as determined by one-way ANOVA in $\delta^{18}\text{O}$ ($F_{5, 109} = 16.23$, $p < 0.001$) and $\delta^{13}\text{C}$ ($F_{5, 109} = 8.17$, $p < 0.001$) of *C. fissilis* sites; and in $\delta^{18}\text{O}$ ($F_{2, 27} = 5.82$, $p = 0.008$) and $\delta^{13}\text{C}$ ($F_{2, 27} = 4.56$, $p = 0.02$) of *C. odorata*.

Annual Isotopic Variation in the 10 Most Recent Rings per Tree

The standard deviations for annual samples within trees ranged from 1.0 to 3.0‰ for $\delta^{18}\text{O}$ (with

averages between 22.0 and 25.3‰), and from 0.3 to 1.2‰ for $\delta^{13}\text{C}$ (with averages between -27.3 and -26.0 ‰; Supplementary Material Table S2). Weighted and averaged annual isotopic data were significantly correlated for $\delta^{18}\text{O}$ (Pearson correlation, $r = 0.95$, $t_{14} = 12$, $p < 0.001$) and $\delta^{13}\text{C}$ (Pearson correlation, $r = 0.92$, $t_{14} = 8.56$, $p < 0.001$; Supplementary Material Figure S2). Moreover, both transformations followed similar isotopic patterns as the 10-year bulk samples (Supplementary Material Figure S3). Additionally, we found no significant correlation between tree-ring width and annual isotopic values for $\delta^{13}\text{C}$ (Pearson correlation, $r = -0.09$, $t_{158} = -1.25$, $p = 0.21$) nor for $\delta^{18}\text{O}$ (Pearson correlation, $r = -0.12$, $t_{158} = -1.50$, $p = 0.13$). Therefore, we decided to use the average annual values as additional samples within the 10-year bulk dataset for further analyses. The 10-year bulk isotopic values were clustered by site and species but had large overlaps (Figure 3).

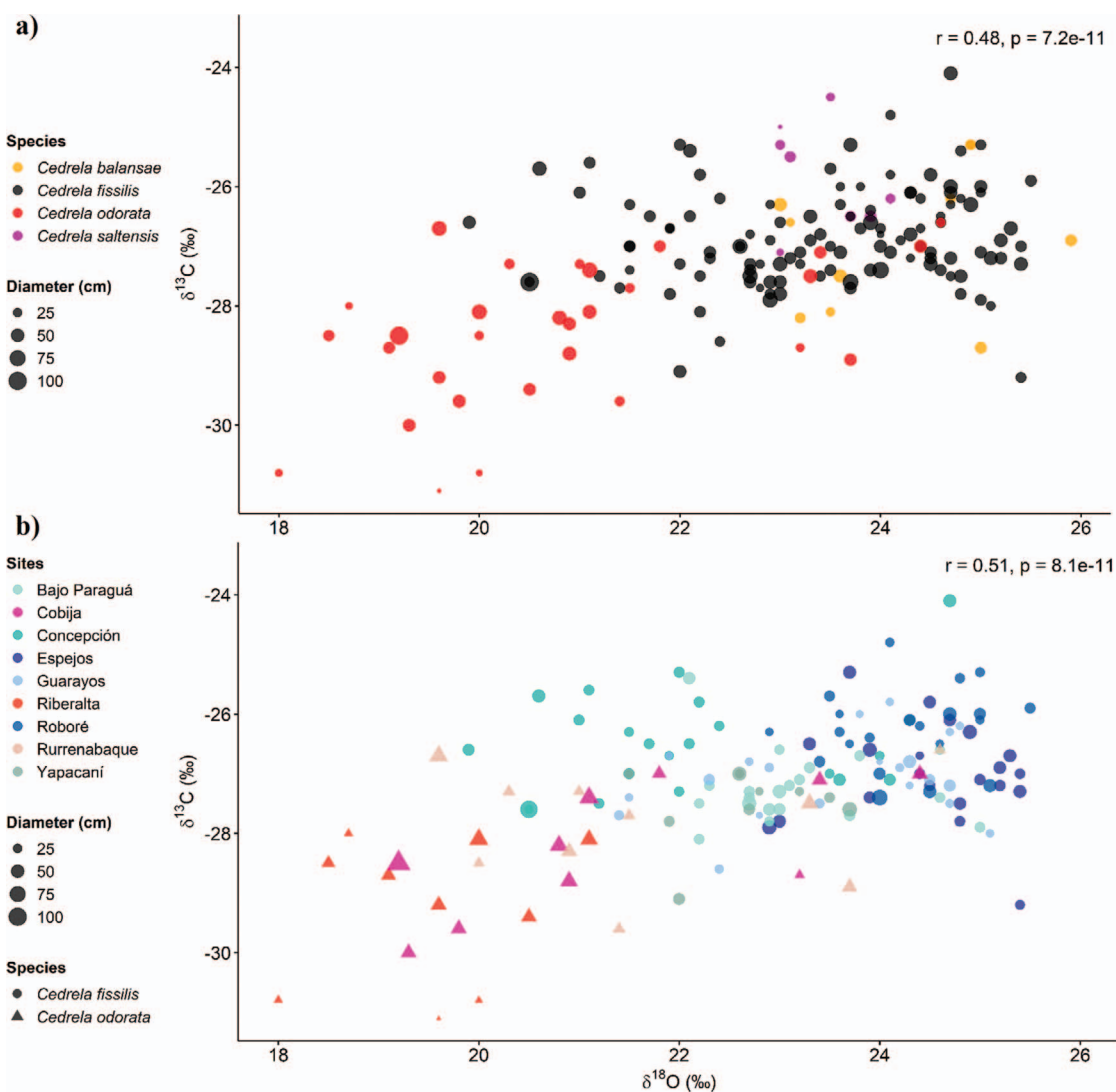


Figure 3. Scatterplots of 10-year bulk samples of $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ stable isotopes of (a) four *Cedrela* species: *C. balansae* (orange circles), *C. fissilis* (black circles), *C. odorata* (red circles), and *C. saltensis* (purple circles); and (b) six *C. fissilis* (circles) and three *C. odorata* (triangles) sample sites.

The correlation between diameter at breast height (DBH) and stable isotopes (Supplementary Material Table S3) was not significant for *C. odorata* ($\delta^{18}\text{O}$ Pearson correlation, $r = -0.003$, $t_{28} = -0.02$, $p = 0.99$; $\delta^{13}\text{C}$ Pearson correlation, $r = 0.30$, $t_{28} = 1.69$, $p = 0.10$) or *C. fissilis* ($\delta^{18}\text{O}$ Pearson correlation, $r = -0.02$, $t_{113} = -0.26$, $p = 0.79$; $\delta^{13}\text{C}$ Pearson correlation, $r = -0.06$, $t_{113} = -0.62$, $p = 0.53$). This was also consistent within each site, with only Cobija, Riberalta and Roboré showing weak relationships between DBH and isotopes (Supplemen-

tary Material Table S3). Overall, any possible effect that size could have on isotopic ratios (Brienen *et al.* 2017) was hardly observed in these patterns.

Relationship Between $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ and Environmental Variables

Isotopic ($\delta^{13}\text{C}$ and $\delta^{18}\text{O}$) composition in *C. odorata* showed no significant correlations with environmental conditions (mean site precipitation or elevation) in the 10-year bulk data. The relationship between stable isotopes in *C.*

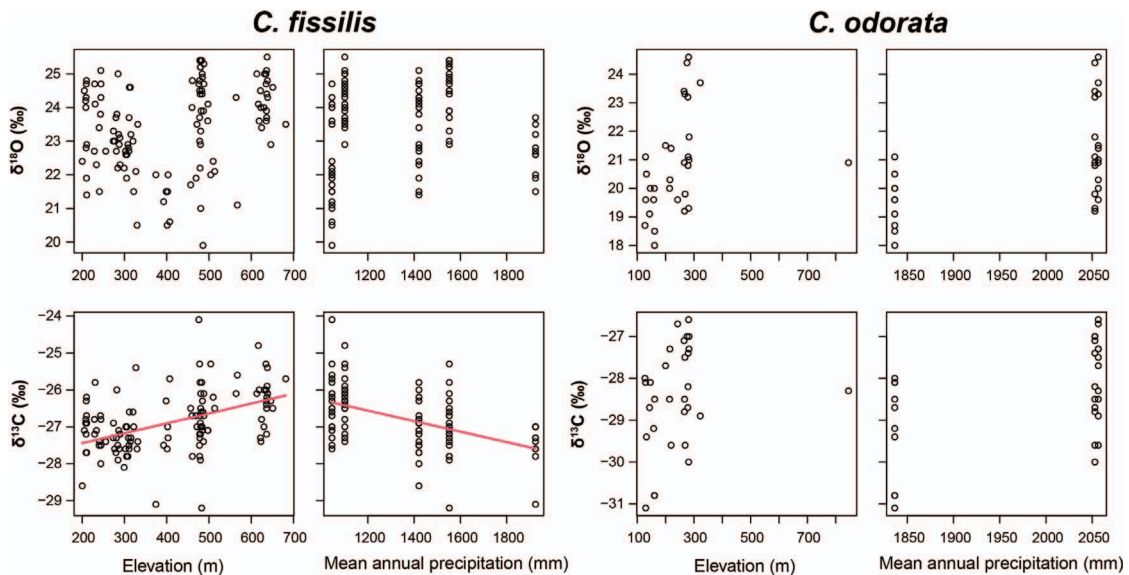


Figure 4. Mixed-effect models to assess the relationship between elevation, mean annual precipitation and stable isotopes ($\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ from 10-year bulk samples). Only significant relationships within the mixed models are indicated (red trend lines).

fissilis and mean precipitation and elevation varied depending on the spatial and temporal scale. At a larger temporal scale, *i.e.* when using 10-year bulk pooled data, we found no significant correlations between $\delta^{18}\text{O}$ in *C. fissilis* and either precipitation or elevation, but $\delta^{13}\text{C}$ had significant correlations with both variables (Table 2, Figure 4). Within-site linear model trends between elevation and isotopic ($\delta^{13}\text{C}$ and $\delta^{18}\text{O}$) composition were not always significant and sometimes not consistent (Table 2, Supplementary Material Figures S1, S4, S5). It is important to note that elevation and precipitation values were highly correlated, and thus their significance when both variables were analyzed together in the model should be interpreted carefully. The variance inflation factor (VIF) of both variables in the models, however, were lower than the usually considered threshold of 5 (1.377 for 10-year bulk, *C. fissilis* and 1.424 for *C. odorata*), which suggested weak collinearity, and thus we decided to retain both variables and show the estimates of these models separately.

Kernel Discriminant Analysis (KDA) for 10-Year-Bulk Samples per Tree for Site Identification

The classification matrix based on *C. fissilis*' 10-year bulk samples per tree (Table 3) estimated an

overall site classification error of 60.2%. Yapacani was the site with the least discriminating isotopic composition, with only 1% of samples correctly assigned to their corresponding origin and most of them wrongly assigned to Bajo Paraguá (82.5%). On the other hand, samples from Bajo Paraguá were mostly correctly identified (61.3%), and those misidentified were predominantly assigned to Espejos (11.8%). Roboré showed the highest classification accuracy (64.8%).

For *C. odorata*, mean total classification error was 70%. Riberalta samples were correctly assigned in 64.6% of cases, with some confusion with Cobija or Rurrenabaque (28.1–7.3%, Table 3). By contrast, the latter two sites showed correct assignments in less than 28% (Rurrenabaque) and 8% (Cobija) of samples.

Species Identification Based on Stable Isotopes

There were statistically significant differences between species group means as determined by one-way ANOVA in $\delta^{18}\text{O}$ ($F_{3,159} = 32.05$, $p < 0.001$) and $\delta^{13}\text{C}$ ($F_{3,159} = 28.57$, $p < 0.001$). The KDA on the 10-year bulk samples showed clear species clustering, especially between *C. fissilis* and *C. odorata* (Figure 3a). Species discrimination for *C. fissilis* was remarkably accurate (95.3%, Table 4).

Table 2. Mixed-effect models to assess potential effect of environmental variables (annual precipitation and elevation) and diameter on $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ composition in *C. fissilis* and *C. odorata*. Analyses were separated for annual and 10-year bulk samples. Mixed-effect model results for annual isotope data ($\delta^{13}\text{C}$ or $\delta^{18}\text{O}$ with precipitation, sample tree nested in site as random factor).

Species	Isotope	Model	Value	Std.err	p-value	df	R ² _m	R ² _c		
A. Annual samples										
<i>C. fissilis</i>	$\delta^{18}\text{O}$	Intercept	23.029	1.1339	0.000	116	0.01	0.16		
		Precipitation	0.0001	0.0005	0.825	116				
		Diameter	0.019	0.0233	0.438	8				
	$\delta^{13}\text{C}$	Intercept	-26.436	0.3960	0.000	116	0.001	0.12		
		Precipitation	0.00001	0.0002	0.945	116				
		Diameter	-0.002	0.0081	0.781	8				
B. 10-year bulk samples										
<i>C. fissilis</i>	$\delta^{18}\text{O}$	Intercept	23.627	1.868	0.000	88	0.002	0.43		
		Precipitation	-0.00002	0.0013	0.991	3				
		Diameter	-0.004	0.0081	0.618	88				
		Intercept	22.267	0.8895	0.000	107			0.12	0.43
		Elevation	0.003	0.0019	0.102	107				
		Diameter	-0.004	0.0076	0.595	107				
	$\delta^{13}\text{C}$	Intercept	-24.755	0.4239	0.000	88	0.24	0.24		
		Precipitation	-0.001	0.0003	0.014	3				
		Diameter	-0.003	0.0053	0.584	88				
		Intercept	-27.734	0.3416	0.000	107			0.23	0.27
		Elevation	0.003	0.0007	0.000	107				
		Diameter	-0.007	0.0052	0.188	107				
	<i>C. odorata</i>	$\delta^{18}\text{O}$	Intercept	1.808	5.3498	0.738	26	0.33	0.33	
			Precipitation	0.009	0.0028	0.173	1			
			Diameter	-0.014	0.0151	0.371	26			
$\delta^{13}\text{C}$		Intercept	20.9361	1.0356	0.000	25	0.02	0.22		
		Elevation	0.0014	0.0026	0.586	25				
		Diameter	-0.011	0.0168	0.530	25				
		Intercept	-39.279	3.8301	0.000	26			0.29	0.29
		Precipitation	0.005	0.0019	0.231	1				
		Diameter	0.013	0.0108	0.255	26				
$\delta^{13}\text{C}$	Intercept	-29.2287	0.6929	0.000	25	0.09	0.24			
	Elevation	0.0001	0.0018	0.974	25					
	Diameter	0.0185	0.0116	0.123	25					

Discriminating between the other three species was more challenging, with higher mean identification errors of 42.5% for *C. odorata*, 82.7% for *C. saltensis*, and 100% for *C. balansae* (Table 4A). Accuracy did not improve by only discriminating between the species with higher sample size (Table 4B).

DISCUSSION

Stable Isotopes' Potential for Identifying Site of Origin

In this study, we assessed isotopic differentiation in *Cedrela* species and their potential to trace wood geographical origin of *C. odorata* and

C. fissilis. We hypothesized that $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ isotopic composition in *Cedrela* would reflect the spatial variation in environmental (McCarroll and Loader 2004) and topographic (Förstel and Hütten 1983) conditions, and thus could be used as a spatial proxy to identify timber origin. Our analyses show promising results but also highlight multiple challenges. Although stable isotopes showed significant correlations with precipitation and elevation at a large spatial scale (*i.e.* comparing average site conditions across sites), these relations were not always consistent within sites, suggesting that other environmental variables, rather than elevation or precipitation, also influence the variability in wood isotopic composition. This was also supported by our

Table 3. Sites classification matrix for the 10-year bulk samples from (A) *C. fissilis* and (B) *C. odorata* species, with the true classes in rows and the estimated classes in columns. Values indicate the assignment of sample origin averaged over 100 runs and their standard deviation (SD). Bolded diagonal values indicate the average correct assignments of samples. Values to the lower left and upper right of the bolded values indicate the average assignments of misclassified samples.

True (%)	Estimated (%)					
A <i>Cedrela fissilis</i>						
Sites	Bajo Paraguá	Concepción	Espejos	Guarayos	Roboré	Yapacaní
Bajo Paraguá	61.3	10.0	11.8	3.6	9.0	4.4
SD	24.5	15.0	16.3	9.3	14.5	13.6
Concepción	22.3	54.3	3.0	7.0	13.0	0.3
SD	22.1	26.3	8.6	13.3	18.4	2.4
Espejos	10.6	0.3	47.3	9.6	31.4	0.8
SD	16.4	3.3	27.5	18.0	23.5	4.4
Guarayos	28.7	8.0	31.6	2.6	24.6	4.6
SD	23.8	14.4	24.4	10.0	21.9	13.4
Roboré	7.5	4.3	17.9	5.5	64.8	0.2
SD	13.1	9.2	19.9	11.0	25.9	1.7
Yapacaní	82.5	8.2	1.7	6.7	0.0	1.0
SD	27.3	20.5	8.4	18.0	0.0	10.0
B <i>Cedrela odorata</i>						
Sites	Cobija	Riberalta	Rurrenabaque			
Cobija	8.3	39.9	51.8			
SD	27.4	39.6	40.8			
Riberalta	28.1	64.6	7.3			
SD	36.7	37.7	20.0			
Rurrenabaque	54.3	18.1	27.7			
SD	39.8	31.2	36.3			

Note: Slight rounding errors may result in sums that are not quite equal to 100%.

Table 4. Classification matrix from 10-year bulk samples of (A) all studied *Cedrela* and (B) the most abundant species, with the true classes in rows and the estimated classes in columns. Values indicate the assignment of sample origin averaged over 100 runs and their standard deviation (SD). Bolded diagonal values indicate the average correct assignments of samples. Values to the lower left and upper right of the bolded values indicate the average assignments of misclassified samples.

True (%)	Estimated (%)			
A All studied species				
Species	<i>C. balansae</i>	<i>C. fissilis</i>	<i>C. odorata</i>	<i>C. saltensis</i>
<i>C. balansae</i>	0.0	99.8	0.3	0.0
SD	0.0	2.5	2.5	0.0
<i>C. fissilis</i>	1.7	95.3	2.9	0.2
SD	2.5	4.2	3.5	1.0
<i>C. odorata</i>	3.4	39.1	57.5	0.0
SD	8.1	23.2	22.3	0.0
<i>C. saltensis</i>	0.3	82.4	0.0	17.3
SD	3.3	32.6	0.0	32.2
B Most abundant species				
Species	<i>C. fissilis</i>	<i>C. odorata</i>		
<i>C. fissilis</i>	96.5	3.5		
SD	4.1	4.1		
<i>C. odorata</i>	40.9	59.1		
SD	20.1	20.1		

Note: Slight rounding errors may result in sums that are not quite equal to 100%.

analyses of annual data, which showed no significant correlations between annual $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values and precipitation (Table 2). Further analysis on site discrimination indicated a limited isotopic potential for tracing *Cedrela* timber geographic origin (Table 3).

Site classification accuracy was highly variable (between 1 and 64.8% for *C. fissilis* and 8.3 to 64.6% for *C. odorata*). This variation in classification accuracy may be explained by a high climatic overlap between some of the sites (similar precipitation conditions), which would also explain the misidentification between some of the sites. Alternatively, there could be species-specific signals or within-site variability in environmental conditions (e.g. in precipitation, elevation, light availability, or edaphic conditions) that may lead to the large differences in accuracy and overall low discrimination. Our discrimination analysis covered a wide spatial resolution, ranging from 3 m to 501 km for *C. fissilis* and from 85 m to 613 km for *C. odorata* samples. The overall low discrimination accuracy in some of the sites is consistent with what has been found for other tropical timber species based on $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ measurements, such as *Erythrophleum* in Cameroon and the Republic of Congo, where average accuracy was 35% (Vlam *et al.* 2018). Similarly, Vlam *et al.* (2018) reported a high variability in classification accuracy, from 46% up to 99%, at lower spatial scales ranging from 14 to 216 km. This suggests that the potential for implementing accurate and consistently reliable wood tracing based on these stable isotopes is likely site- and species-dependent. Also, our results highlight that the use of $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ alone may be insufficient to reliably account for variability in environmental conditions and scale.

Interannual variation in tree-ring $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ has been related to environmental conditions (McCarroll and Loader 2004; West *et al.* 2006). However, contrary to our expectations and to previous work (Dansgaard 1964; Gonfiantini *et al.* 2001), we found no significant correlations between annual isotopic data and precipitation or elevation (Table 2A). Low correlations between precipitation and $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ have also been reported in other tree species (Poussart and Schrag 2005; Cullen and Grierson 2007; Schollaen *et al.* 2013; Baker *et al.* 2015), and have been attributed to changing

limiting factors of tree growth between years (Helle and Schleser 2004; Paredes-Villanueva *et al.* 2016). For example, because *Cedrela* is a relatively light-sensitive tree (Brienen and Zuidema 2006; Brienen *et al.* 2010), light availability might show a greater effect on tree development than precipitation in some sites (Brienen *et al.* 2010) but not in others. In theory, this periodical shift in the variable limiting growth could be taken into account by using annual isotopic data. However, in our study, using annual variation did not improve discrimination ability (mean error 70.1%), but rather added noise (Supplementary Material Table S4). Another source of uncertainty in our models may be the insufficient resolution and large scale of the environmental data (Kurita *et al.* 2009). Studying isotopes on a small scale while the limiting factor operates at a large scale can affect statistical analysis and consequent discrimination among sites. Indeed, large-scale environmental variation seemed to reflect site characteristics more reliably in our dataset when using 10-year pooled samples. The significant relationship between environmental conditions – such as precipitation and elevation – and the 10-year bulk isotopic signatures found on our exploratory analyses again reinforced the importance of considering the resolution and scale of environmental data. Analyzing longer time series of tree rings per tree could be a path forward to increase signals detection and accuracy of the results.

Although the potential of stable isotopes to trace timber geographical origin has previously been demonstrated in some species (Förstel and Hützen 1983; Boner *et al.* 2007; Kagawa and Leavitt 2010; Förstel *et al.* 2011; Vlam *et al.* 2018; Watkinson *et al.* 2020), timber tracing with a fine spatial resolution (*i.e.* higher precision) remains a challenge. We used a wide range of annual precipitation and elevation regimes to assess if their relation with trees will be mirrored in their isotopic composition ($\delta^{13}\text{C}$ and $\delta^{18}\text{O}$). The overall inconsistencies between these environmental variables and our annual/10-year bulk isotopic data, in addition to the low discrimination among sites, suggest an influence by other governing factors at large spatiotemporal scales. For example, soil properties and El Niño-Southern Oscillation (ENSO), may not only influence isotopic composition in trees directly but also have marked effects depending on species

and habitat. Previous research on $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ in coca (*Erythroxylum coca*) from South America found significant isotopic differences among sample sites and attributed it to differences in soil (for $\delta^{15}\text{N}$) and length of the wet season (for $\delta^{13}\text{C}$) (Ehleringer *et al.* 2000). Soil characteristics, particularly soil water availability and nutrients (Medina *et al.* 1995; Oliveira-Filho *et al.* 1998), have also been shown to have an impact on tree growth in dry forests (Toledo *et al.* 2011), and it is therefore likely to influence wood isotopic composition. This impact shows a spatial gradient (Murphy and Lugo 1986; Cecon *et al.* 2006) that progressively shifts in wetter forests where light variation is more important for growth (Engelbrecht *et al.* 2007; Brienen *et al.* 2010). However, where soil chemicals are a limiting factor driving tree growth, it is important to further study which chemicals leave a signal in their wood composition.

The temporal-scale influence of ENSO is prevalent across South American forests, including on the isotopic composition of *Cedrela* species in Bolivia (Vuille and Werner 2005; Brienen *et al.* 2012; Baker *et al.* 2015). Vuille and Werner (2005) found higher $\delta^{18}\text{O}$ values during the El Niño phase (drier) of ENSO, and lower $\delta^{18}\text{O}$ values during La Niña events. Furthermore, at small spatial scale, precipitation was found to be associated with local circulation, and at a larger spatial scale, isotopic variations of precipitation were found to depend on the rain-out process in the surrounding region (Kurita *et al.* 2009). Although the rain-out history at a regional scale may be reflected in isotopic signal of precipitation, how stable isotopes composition varies in trees will depend on the site- and species-specific responses to these large-scale environmental factors (Helle and Schleser 2004; Poussart and Schrag 2005; van der Sleen *et al.* 2017). To our knowledge, temporal variability of stable isotopes has been studied for different species and sites in Bolivia (Brienen *et al.* 2012; Nijmeijer 2012; van der Sleen 2014; Baker *et al.* 2015; van der Sleen *et al.* 2015). However, for timber tracing, a good spatial representation is required. Although stable isotopes show promise, a better understanding of the between-species and within-sites variability is required before discrimination analysis can be widely and reliably implemented.

Stable Isotopes to Identify *Cedrela* Species

Our results showed distinct value ranges for each of the species' isotopic composition, with higher values for *C. fissilis* (ca. 24‰ $\delta^{18}\text{O}$, ca. -27‰ $\delta^{13}\text{C}$) compared to *C. odorata* (ca. 20‰ $\delta^{18}\text{O}$, ca. -29‰ $\delta^{13}\text{C}$) (Figure 3). ^{13}C discrimination is a function of the ratio of internal to ambient CO_2 concentrations ($C_i:C_a$) and water-use efficiency (Farquhar *et al.* 1982; Farquhar and Richards 1984). Under humid conditions $C_i:C_a$ is larger resulting in more discrimination against ^{13}C (more negative $\delta^{13}\text{C}$ values). Under dry conditions discrimination is less, resulting in more positive $\delta^{13}\text{C}$ values because of a relative enrichment in ^{13}C . This was in line with our results, where the $\delta^{13}\text{C}$ 10-year bulk values ranged from -31‰ for the moist-adapted species (*C. odorata*) to -24‰ for the dry-tolerant species (*C. fissilis*).

Although stable isotopes have been previously implemented to identify animal species (Oliveira *et al.* 2011; Chouvelon *et al.* 2014; Militão *et al.* 2014; Hamer *et al.* 2015; Ditttrich *et al.* 2017), our study represents a first exploration on their potential use for taxonomic and forensics purposes in timber species. In our case, using 10-year bulk samples seemed to render highly accurate identification for *C. fissilis* with up to 95.3% accuracy, but seemed insufficient to reliably identify other *Cedrela* species, particularly *C. balansae*, which showed 100% error. However, the worst performing species in our classification analyses were also the least replicated ones, indicating the need for larger sample sizes for each species in future work. Consistent with our results, previous discrimination analyses using Direct Analysis in Real Time (DART) coupled to Time-of-Flight Mass Spectrometry (TOFMS) also found *C. balansae* challenging to identify, with the highest error of 46.1% compared to *C. fissilis* which presented the highest accuracy (91.3%) (Paredes-Villanueva *et al.* 2018). The current data, though limited, suggest that identifying *C. balansae* via isotopic signatures may require their use in combination with other (*e.g.* genetics, anatomical, Near Infrared Spectroscopy) approaches.

An additional source of identification error that should be taken into account concerns updates on the species classifications based on herbarium

collections. Rapidly advancing tropical botanical studies mean that the morphological characteristics to tease apart *Cedrela* species are constantly being updated, which makes accurate identification via botanical samples sometimes challenging (Zapater *et al.* 2004; Koecke *et al.* 2015; Palacios *et al.* 2019). Species classification based on both morphological and genetic properties might help us to perform chemical discrimination for then well-established species. In addition, it will provide insights into the underlying causes of identification errors. These protocols can consequently be fine-tuned to discriminate between problematic species.

How much isotopic variation is there among all traded timber species? Are there any other species having the same isotopic values as any specific *Cedrela* species, which could increase the error rate for identification? Further work is needed to map the expected isotopic range for more timber species, in order to contextualize our results and provide widely applicable tools for sustainable timber trading.

Considerations for ^{13}C and ^{18}O Stable Isotope Applications in Timber Tracing

Some considerations have to be taken into account to interpret our ^{13}C and ^{18}O results. First, a light-demanding species may differ in its isotopic composition if samples are collected at disturbed areas or if wood tissue was formed during clearing events (Brienen *et al.* 2010; van der Sleen *et al.* 2014). In our case, samples were collected in areas where there was previous intervention or use, as well as in dense forests with closed canopy. However, tree sensitivity and its isotopic composition may also be influenced by temporally shifting limiting factors (Helle and Schleser 2004) at a specific period. In case there is a possibility that the intervention in a forest or light availability plays an important role in the tree isotopic composition, we recommend taking into account the intervention degree in future discrimination analyses.

Second, we expected isotopic values to be reliable proxies for climate data (*e.g.* precipitation). Indeed, *Cedrela* trees' shallow roots would suggest a strong correlation between $\delta^{18}\text{O}$ in trees and precipitation (Brienen *et al.* 2012). However, in our study, precipitation did not show high variation among

C. fissilis sites (Table 1) nor significant correlation with the $\delta^{18}\text{O}$ (Table 2, Figure 4). This suggests that precipitation may not always be the main driving factor of $\delta^{18}\text{O}$ composition in *Cedrela* wood. Other possible oxygen sources that can have an effect on isotopic variation in plants include humidity, soil moisture, groundwater (Aggarwal *et al.* 2004), and clouds (Anchukaitis *et al.* 2008; Anchukaitis and Evans 2010).

Third, isotopic variation also depends on tree age and size (ontogenetic development phase) (Broadmeadow *et al.* 1992; McCarroll and Loader 2004; van der Sleen *et al.* 2015; Brienen *et al.* 2016). However, we found no supporting evidence for this effect influencing the discrimination potential of our isotope samples. Tree size (DBH) did not have a significant effect in the isotopic composition of our trees. We also found no significant correlation between DBH and stable isotopes in *C. odorata*, nor in *C. fissilis*. Similarly, removing trees with exceptionally small or large DBH (*i.e.* 2 *C. odorata* samples with <15 and >70 cm, and 4 *C. fissilis* samples with <20 and >70 cm) to minimize this potential size/age effects, had little effect on mean discrimination accuracy (+1.2% in *C. fissilis* and -0.5% in *C. odorata*).

In addition, during an analysis of timber with unknown origin, estimating the possible period of tree harvesting will be a minimum requirement to correct for the overall decrease in $\delta^{13}\text{C}$ in tree biomass since the onset of the Industrial Revolution period (McCarroll and Loader 2004; van der Sleen *et al.* 2017). Finally, the low spatial variation in $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ could be improved by assessing multiple tree-isotope proxies or combining them with rare earth and trace elements (English *et al.* 2001; Kelly *et al.* 2005; Joebstl *et al.* 2010) to increase discrimination among sites.

CONCLUSIONS

Cedrela species are among the most important traded timbers species and have suffered from over-exploitation. We aimed at assessing whether timber isotopic composition (based on $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$) can be used to differentiate *Cedrela* species and sites of origin, providing a key regulatory tool for tropical forests. This approach could potentially improve the verification of timber origin certificates in

accordance with regulations. However, our analyses reveal limited potential for discrimination of geographical origin using wood $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$. Sites assignment accuracy was low, at 30% for *C. odorata* and 39.8% for *C. fissilis*. This low geographical discrimination potential, using carbon and oxygen isotopic imprints alone, may result from multiple factors, including high climatic overlap between certain sites, high within-site variability, and lack of high-resolution environmental data. In addition, using precipitation and elevation as proxies to infer site of origin at a fine resolution (specific tree location) remains a challenge, given the high variability of isotopic composition ($\delta^{13}\text{C}$ and $\delta^{18}\text{O}$) within trees. Yet, our results suggest that it is possible to distinguish between species based on stable isotopes. In particular, *C. fissilis* can be identified with a remarkably high accuracy (95.3%). We conclude that although much work is still needed to be able to use stable isotopes as a sole tool to identify tropical trees provenances, this methodology already shows potential for discrimination of some important timber species. Further developments in this direction will contribute to improve timber origin classification and thereby may become a tool in the international fight against illegal trade of (precious) tropical timbers.

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REFERENCES CITED

- ABT, 2017. *Audiencia Pública Inicial y Parcial de Rendición de Cuentas (de Enero Hasta Agosto, Gestión 2017)* [PowerPoint slides]. Autoridad de Fiscalización y Control Social de Bosques y Tierra (ABT), Santa Cruz, Bolivia. <http://www.abt.gob.bo/images/stories/audienciaPublica/2017/audiencia-inicial-parcial/AUDIENCIA%20PUBLICA%20INICIAL-PARCIAL%202017%20ABT.pdf>.
- Abt Associates Inc., 2006. *Illegal Logging: A Market-based Analysis of Trafficking in Illegal Timber*. Cambridge, MA; 56 pp.
- Aggarwal, P. K., K. Fröhlich, K. M. Kulkarni, and L. L. Gourcy, 2004. Stable isotope evidence for moisture sources in the Asian Summer Monsoon under present and past climate regimes. *Geophysical Research Letters* 31:1–4.
- Anchukaitis, K. J., and M. N. Evans, 2010. Tropical cloud forest climate variability and the demise of the Monteverde golden toad. *Proceedings of the National Academy of Sciences* 107:5036–5040.
- Anchukaitis, K. J., M. N. Evans, N. T. Wheelwright, and D. P. Schrag, 2008. Stable isotope chronology and climate signal calibration in neotropical montane cloud forest trees. *Journal of Geophysical Research: Biogeosciences* 113, doi:10.1029/2007JG000613.
- Baker, J. C. A., S. F. P. Hunt, S. J. Clerici, R. J. Newton, S. H. Bottrell, M. J. Leng, T. H. E. Heaton, G. Helle, J. Argollo, M. Gloor, and R. J. W. Brienen, 2015. Oxygen isotopes in tree rings show good coherence between species and sites in Bolivia. *Global and Planetary Change* 133:298–308.
- Baudat, G., and F. Anouar, 2000. Generalized discriminant analysis using a kernel approach. *Neural Computation* 12:2385–2404.
- Blaser, J., 2010. *Forest Law Compliance and Governance in Tropical Countries: A Region-by-region Assessment of the Status of Forest Law Compliance and Governance in the Tropics, and Recommendations for Improvement*. FAO/ITTO; 27 pp.
- Boner, M., T. Sommer, C. Erven, and H. Förstel, 2007. Stable isotopes as a tool to trace back the origin of wood. In *Proceedings of the International Workshop "Fingerprinting Methods for the Identification of Timber Origins"*, edited by Degen, B., Special Issue 321, pp. 3–5. Johann Heinrich von Thünen-Institut, Braunschweig, Germany.
- Booth, A. L., M. J. Wooller, T. Howe, and N. Haubstock, 2010. Tracing geographic and temporal trafficking patterns for marijuana in Alaska using stable isotopes (C, N, O and H). *Forensic Science International* 202:45–53.
- Bowen, G. J., and J. Revenaugh, 2003. Interpolating the isotopic composition of modern meteoric precipitation. *Water Resources Research* 39:1299, doi.org/10.1029/2003WR002086
- Brauning, A., F. Volland-Voigt, I. Burchardt, O. Ganzhi, T. Naus, and T. Peters, 2009. Climatic control of radial growth of *Cedrela montana* in a humid mountain rainforest in southern Ecuador. *Erdkunde* 63:337–345.
- Brienen, R. J. W., E. Gloor, S. Clerici, R. Newton, L. Arppe, A. Boom, S. Bottrell, M. Callaghan, T. Heaton, S. Helama, G. Helle, M. J. Leng, K. Mielikäinen, M. Oinonen, and M. Timonen, 2017. Tree height strongly affects estimates of water-use efficiency responses to climate

- and CO₂ using isotopes. *Nature Communications* 8:1–10. <https://doi.org/10.1038/s41467-017-00225-z>.
- Brienen, R. J. W., G. Helle, T. L. Pons, J.-L. Guyot, and M. Gloor, 2012. Oxygen isotopes in tree rings are a good proxy for Amazon precipitation and El Niño-Southern Oscillation variability. *Proceedings of the National Academy of Sciences* 109: 16957–16962.
- Brienen, R. J. W., J. Schöngart, and P. A. Zuidema, 2016. Tree rings in the tropics: Insights into the ecology and climate sensitivity of tropical trees. In *Tropical Tree Physiology: Adaptations and Responses in a Changing Environment*, edited by Goldstein, G., and L. Santiago, pp. 439–461. Springer, Cham.
- Brienen, R. J. W., and P. A. Zuidema, 2005. Relating tree growth to rainfall in Bolivian rain forests: A test for six species using tree ring analysis. *Oecologia* 146:1–12.
- Brienen, R. J. W., and P. A. Zuidema, 2006. Lifetime growth patterns and ages of Bolivian rain forest trees obtained by tree ring analysis. *Journal of Ecology* 94:481–493.
- Brienen, R. J. W., P. A. Zuidema, and M. Martínez-Ramos, 2010. Attaining the canopy in dry and moist tropical forests: Strong differences in tree growth trajectories reflect variation in growing conditions. *Oecologia* 163:485–496.
- Broadmeadow, M. S. J., H. Griffiths, C. Maxwell, and A. M. Borland, 1992. The carbon isotope ratio of plant organic material reflects temporal and spatial variations in CO₂ within tropical forest formations in Trinidad. *Oecologia* 89:435–441.
- Bunn, A., M. Korpela, F. Biondi, F. Campelo, P. Mérian, F. Qeadan, and C. Zang, 2021. dplR: Dendrochronology Program Library in R. R package version 1.7.2 2022.
- Ceccon, E., P. Huante, and E. Rincón, 2006. Abiotic factors influencing tropical dry forests regeneration. *Brazilian Archives of Biology and Technology* 49:305–312.
- Chouvelon, T., F. Caurant, Y. Cherel, B. Simon-Bouhet, J. Spitz, and P. Bustamante, 2014. Species- and size-related patterns in stable isotopes and mercury concentrations in fish help refine marine ecosystem indicators and provide evidence for distinct management units for hake in the Northeast Atlantic. *ICES Journal of Marine Science* 71:1073–1087.
- CITES, 2019. Proposals for Amendment of Appendices I and II - Eighteenth Meeting of the Conference of the Parties 2019. <https://cites.org/eng/cop/18/prop/index.php>
- Cullen, L. E., and P. F. Grierson, 2007. A stable oxygen, but not carbon, isotope chronology of *Callitris columellaris* reflects recent climate change in north-western Australia. *Climatic Change* 85:213–229.
- Dansgaard, W., 1964. Stable isotopes in precipitation. *Tellus* 16:436–468.
- Degen, B., and M. Fladung, 2007. Use of DNA-markers for tracing illegal logging. In *Proceedings of the International Workshop "Fingerprinting Methods for the Identification of Timber Origins"*, edited by Degen, B., Special Issue 321, pp. 6–14. Johann Heinrich von Thünen-Institut, Braunschweig, Germany.
- Dittrich, C., U. Struck, and M.-O. Rödel, 2017. Stable isotope analyses - A method to distinguish intensively farmed from wild frogs. *Ecology and Evolution* 7:2525–2534.
- Driscoll, A. W., J. D. Howa, N. Q. Bitter, and J. R. Ehleringer, 2020. A predictive spatial model for roasted coffee using oxygen isotopes of α -cellulose. *Rapid Communications in Mass Spectrometry* 34:e8626.
- Dünisch, O., J. Bauch, and L. Gasparotto, 2002. Formation of increment zones and intraannual growth dynamics in the xylem of *Swietenia macrophylla*, *Carapa guianensis*, and *Cedrela odorata* (Meliaceae). *IAWA Journal* 23:101–119.
- Duong, T., 2007. ks: Kernel density estimation and kernel discriminant analysis for multivariate data in R. *Journal of Statistical Software* 21:1–16.
- Duong, T., 2017. ks: Kernel Smoothing. R package version 1.10.6 <https://cran.r-project.org/web/packages/ks/>
- Ehleringer, J. R., G. J. Bowen, L. A. Chesson, A. G. West, D. W. Podlesak, and T. E. Cerling, 2008. Hydrogen and oxygen isotope ratios in human hair are related to geography. *Proceedings of the National Academy of Sciences* 105:2788–2793.
- Ehleringer, J. R., J. F. Casale, M. J. Lott, and V. L. Ford, 2000. Tracing the geographical origin of cocaine: Cocaine carries a chemical fingerprint from the region where the coca was grown. *Nature* 408:311–312.
- Engelbrecht, B. M. J., L. S. Comita, R. Condit, T. A. Kursar, M. T. Tyree, B. L. Turner, and S. P. Hubbell, 2007. Drought sensitivity shapes species distribution patterns in tropical forests. *Nature* 447:80–82.
- English, N. B., J. L. Betancourt, J. S. Dean, and J. Quade, 2001. Strontium isotopes reveal distant sources of architectural timber in Chaco Canyon, New Mexico. *Proceedings of the National Academy of Sciences* 98:11891–11896.
- Evans, M. N., and D. P. Schrag, 2004. A stable isotope-based approach to tropical dendroclimatology. *Geochimica et Cosmochimica Acta* 68:3295–3305.
- Farquhar, G. D., M. H. O’Leary, and J. A. Berry, 1982. On the relationship between carbon isotope discrimination and the intercellular carbon dioxide concentration in leaves. *Functional Plant Biology* 9:121–137.
- Farquhar, G. D., and R. A. Richards, 1984. Isotopic composition of plant carbon correlates with water-use efficiency of wheat genotypes. *Australian Journal of Plant Physiology* 11: 539–552.
- Förstel, H., M. Boner, A. Höltnen, M. Fladung, B. Degen, and J. Zahnen, 2011. *Fighting Illegal Logging through the Introduction of a Combination of the Isotope Method for Identifying the Origins of Timber and DNA Analysis for Differentiation of Tree Species*. WWF Germany, Berlin, Germany.
- Förstel, H., and H. Hützen, 1983. Oxygen isotope ratios in German groundwater. *Nature* 304:614–616.
- Gonfiantini, R., M.-A. Roche, J.-C. Olivry, J.-C. Fontes, and G. M. Zuppi, 2001. The altitude effect on the isotopic composition of tropical rains. *Chemical Geology* 181:147–167.
- Gorlova, E. N., O. A. Krylovich, A. V. Tiunov, B. F. Khasanov, D. D. Vasyukov, and A. B. Savinetsky, 2015. Stable-isotope analysis as a method of taxonomical identification of archaeozoological material. *Archaeology, Ethnology and Anthropology of Eurasia* 43:110–121.
- Hamer, S., A. Weghorst, L. Auckland, E. Roark, O. Strey, P. Teel, and G. Hamer, 2015. Comparison of DNA and carbon and nitrogen stable isotope-based techniques for identification of prior vertebrate hosts of ticks. *Journal of Medical Entomology* 52:1043–1049.

- Helle, G., and G. H. Schleser, 2004. Interpreting climate proxies from tree-rings. In *The Climate in Historical Times: Towards a Synthesis of Holocene Proxy Data and Climate Models*, edited by Fischer, H., T. Kumke, G. Lohmann, G. Flöser, H. Miller, H. von Storch, and J. F. W. Negendank, pp. 129–148. Springer, Berlin, Heidelberg.
- Hobson, K. A., 1999. Tracing origins and migration of wildlife using stable isotopes: A review. *Oecologia* 120:314–326.
- Hobson, K. A., and L. I. Wassenaar, 1999. Stable isotope ecology: An introduction. *Oecologia* 120:312–313.
- Jenkins, H. S., 2009. *Amazon Climate Reconstruction Using Growth Rates and Stable Isotopes of Tree Ring Cellulose from the Madre de Dios Basin, Peru*. Ph.D. dissertation, Duke University.
- Jianbang G., P. O. Cerutti, M. Masiero, D. Pettenella, N. Andrighetto, and T. Dawson, 2016. *Quantifying Illegal Logging and Related Timber Trade*. International Union of Forest Research Organizations (IUFRO), Vienna.
- Joebstl, D., D. Bandoniene, T. Meisel, and S. Chatzistathis, 2010. Identification of the geographical origin of pumpkin seed oil by the use of rare earth elements and discriminant analysis. *Food Chemistry* 123:1303–1309.
- Kagawa, A., and S. W. Leavitt, 2010. Stable carbon isotopes of tree rings as a tool to pinpoint the geographic origin of timber. *Journal of Wood Science* 56:175–183.
- Kassambara, A., 2017. ggpubr: 'ggplot2' Based Publication Ready Plots. R package version 0.1.6. <http://www.sthda.com/english/rpkgs/ggpubr>.
- Kelly, S., K. Heaton, and J. Hoogewerff, 2005. Tracing the geographical origin of food: The application of multi-element and multi-isotope analysis. *Trends in Food Science and Technology* 16:555–567.
- Koecke, A. V., A. N. Muellner-Riehl, O. M. Cáceres, and T. D. Pennington, 2015. *Cedrela ngobe* (Meliaceae), a new species from Panamá and Costa Rica. *Edinburgh Journal of Botany* 72:225–233.
- Körner, C., 1998. A re-assessment of high elevation treeline positions and their explanation. *Oecologia* 115:445–459.
- Körner, C., G. D. Farquhar, and Z. Roksandic, 1988. A global survey of carbon isotope discrimination in plants from high altitude. *Oecologia* 74:623–632.
- Kurita, N., K. Ichiyanagi, J. Matsumoto, M. D. Yamanaka, and T. Ohata, 2009. The relationship between the isotopic content of precipitation and the precipitation amount in tropical regions. *Journal of Geochemical Exploration* 102: 113–122.
- Lowe, A. J., E. E. Dormontt, M. J. Bowie, B. Degen, S. Gardner, D. Thomas, C. Clarke, A. Rimbawanto, A. Wiedenhoft, Y. Yin, and N. Sasaki, 2016. Opportunities for improved transparency in the timber trade through scientific verification. *BioScience* 66:990–998.
- McCarroll, D., and N. J. Loader, 2004. Stable isotopes in tree rings. *Quaternary Science Reviews* 23:771–801.
- Medina, E., H. A. Mooney, and S. H. Bullock, 1995. *Seasonally Dry Tropical Forests*. Cambridge University Press, Cambridge.
- Militão, T., E. Gómez-Díaz, A. Kaliontzopoulou, and J. González-Solis, 2014. Comparing multiple criteria for species identification in two recently diverged seabirds. *PLOS ONE* 9:e115650.
- Mostacedo, B., and T. Fredericksen, 1999. Regeneration status of important tropical forest tree species in Bolivia: Assessment and recommendations. *Forest Ecology and Management* 124:263–273.
- Mostacedo, B., and T. Fredericksen, 2001. *Regeneración y Silvicultura de Bosques Tropicales en Bolivia*. Editora El País, Santa Cruz, Bolivia.
- Mostacedo, B., J. Justiniano, M. Toledo, and T. Fredericksen, 2003. *Guía Dendrológica de Especies Forestales de Bolivia*. Proyecto de Manejo Forestal Sostenible (BOLFOR), Santa Cruz, Bolivia.
- Murphy, P. G., and A. E. Lugo, 1986. Ecology of tropical dry forest. *Annual Review of Ecology and Systematics* 17: 67–88.
- NASA/METI/AIST/Japan Spacesystems, and U.S./Japan ASTER Science Team, 2019. *ASTER Global Digital Elevation Model V003 [Data set]*. NASA EOSDIS Land Processes DAAC. <https://doi.org/10.5067/ASTER/ASTGTM.003>.
- Navarro-Cerrillo, R. M., N. Agote, F. Pizarro, C. J. Ceacero, and G. Palacios, 2013. Elements for a non-detriment finding of *Cedrela* spp. in Bolivia — A CITES implementation case study. *Journal for Nature Conservation* 21:241–252.
- Navarro, G., 2011. *Clasificación de la Vegetación de Bolivia*. Centro de Ecología Difusión Simón I. Patiño, Santa Cruz, Bolivia.
- Nijmeijer, A., 2012. *Physiological Responses of a Tropical Tree to Elevated CO₂: A Century Long Evaluation of Pseudolmedia laevis Trees Using Stable Carbon Isotope Values from Tree Rings*. Ph.D. dissertation, Wageningen University, Wageningen, the Netherlands.
- Oliveira-Filho, A. T., N. Curi, E. A. Vilela, and D. A. Carvalho, 1998. Effects of canopy gaps, topography, and soils on the distribution of woody species in a central Brazilian deciduous dry forest. *Biotropica* 30:362–375.
- Oliveira, E. J. V. M., L. S. Sant'Ana, C. Ducatti, J. C. Denadai, and C. R. de Souza Kruliski, 2011. The use of stable isotopes for authentication of gadoid fish species. *European Food Research and Technology* 232:97–101.
- Ometto, J. P. H. B., L. B. Flanagan, L. A. Martinelli, M. Z. Moreira, N. Higuchi, and J. R. Ehleringer, 2002. Carbon isotope discrimination in forest and pasture ecosystems of the Amazon Basin, Brazil. *Global Biogeochemical Cycles* 16: 56-51–56-10.
- Orvis, K. H., and H. D. Grissino-Mayer, 2002. Standardizing the reporting of abrasive papers used to surface tree-ring samples. *Tree-Ring Research* 58:47–50.
- Oulhote, Y., B. Le Bot, S. Deguen, and P. Glorennec, 2011. Using and interpreting isotope data for source identification. *Trends in Analytical Chemistry* 30:302–312.
- Palacios, W. A., J. Santiana, and J. Iglesias, 2019. A new species of *Cedrela* (Meliaceae) from the eastern flanks of Ecuador. *Phytotaxa* 393:84–88.
- Paredes-Villanueva, K., E. Espinoza, J. Ottenburghs, M. G. Sterken, F. Bongers, and P. A. Zuidema, 2018. Chemical differentiation of Bolivian *Cedrela* species as a tool to trace illegal timber trade. *Forestry: An International Journal of Forest Research* 91:603–613.

- Paredes-Villanueva, K., L. López, and R. M. Navarro-Cerrillo, 2016. Regional chronologies of *Cedrela fissilis* and *Cedrela angustifolia* in three forest types and their relation to climate. *Trees - Structure and Function* 30:1581–1593.
- Pinheiro, J., D. Bates, and R-core, 2020. *nlme: Linear and Non-linear Mixed Effects Models*. R package version 3.1–148 2020, <https://CRAN.R-project.org/package=nlme>.
- Poussart, P. F., and D. P. Schrag, 2005. Seasonally resolved stable isotope chronologies from northern Thailand deciduous trees. *Earth and Planetary Science Letters* 235:752–765.
- R Development Core Team, 2017. *R: A Language and Environment for Statistical Computing*. R Foundation for Statistical Computing, Vienna, Austria
- Richter, H. G., and M. J. Dallwitz, 2000. *Commercial Timbers: Descriptions, Illustrations, Identification, and Information Retrieval. In English, French, German, Portuguese, and Spanish. Version: 25th June 2009*. <http://delta-intkey.com>.
- Rossmann, A., 2001. Determination of stable isotope ratios in food analysis. *Food Reviews International* 17:347–381.
- Rozanski, K., L. Araguás-Araguás, and R. Gonfiantini, 1993. Isotopic patterns in modern global precipitation. In *Climate Change in Continental Isotopic Records*, edited by P. K. Swart, K. C. Lohmann, J. Mckenzie, and S. Savin, pp. 1–36. Geophysical Monograph Series, Volume 78. American Geophysical Union, Washington, D.C.
- Rubenstein, D. R., and K. A. Hobson, 2004. From birds to butterflies: Animal movement patterns and stable isotopes. *Trends in Ecology and Evolution* 19:256–263.
- Schollaen, K., I. Heinrich, B. Neuwirth, P. J. Krusic, R. D. D'Arrigo, O. Karyanto, and G. Helle, 2013. Multiple tree-ring chronologies (ring width, $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$) reveal dry and rainy season signals of rainfall in Indonesia. *Quaternary Science Reviews* 73:170–181.
- Schulman, E., 1956. *Dendroclimatic Changes in Semiarid America*. University of Arizona Press, Tucson, Arizona.
- SENAMHI, 2018. Servicio Nacional de Meteorología e Hidrología, Bolivia 2018. <http://www.senamhi.gob.bo/>.
- Toledo, M., L. Poorter, M. Peña-Claros, A. Alarcón, J. Balcázar, C. Leño, J. C. Licona, O. Llanque, V. Vroomans, and P. Zuidema, 2011. Climate is a stronger driver of tree and forest growth rates than soil and disturbance. *Journal of Ecology* 99:254–264.
- University of East Anglia Climatic Research Unit, 2008. *Climatic Research Unit (CRU): Time-Series (TS) Datasets of Variations in Climate with Variations in Other Phenomena v3*. NCAS British Atmospheric Data Centre. <http://catalogue.ceda.ac.uk/uuid/3f894480cc48e1cbc29a5ee12d8542d>
- van der Sleen, J. P., 2014. *Environmental and Physiological Drivers of Tree Growth: A Pan-tropical Study of Stable Isotopes in Tree Rings*. Ph.D. dissertation, Wageningen University, Wageningen, the Netherlands.
- van der Sleen, J. P., P. Groenendijk, M. Vlam, N. P. Anten, A. Boom, F. Bongers, T. L. Pons, G. Terburg, and P. A. Zuidema, 2015. No growth stimulation of tropical trees by 150 years of CO_2 fertilization but water-use efficiency increased. *Nature Geoscience* 8:24–28.
- van der Sleen, J. P., P. A. Zuidema, and T. L. Pons, 2017. Stable isotopes in tropical tree rings: Theory, methods and applications. *Functional Ecology* 31:1674–1689.
- van der Sleen, J. P., P. A. Zuidema, C. C. Soliz-Gamboa, G. Helle, T. L. Pons, and N. P. R. Anten, 2014. Understanding causes of tree growth response to gap formation: $\delta^{13}\text{C}$ -values in tree rings reveal a predominant effect of light. *Trees - Structure and Function* 28:439–448.
- Versini, G., A. Monetti, and F. Reniero, 1997. Monitoring authenticity and regional origin of wines by natural stable isotope ratios analysis. In *Wine: Nutritional and Therapeutic Benefits*, edited by Watkins, T. R. American Chemical Society, Washington, D.C.
- Vlam, M., G. A. de Groot, A. Boom, P. Copini, I. Laros, K. Veldhuijzen, D. Zakamdi, and P. A. Zuidema, 2018. Developing forensic tools for an African timber: Regional origin is revealed by genetic characteristics, but not by isotopic signature. *Biological Conservation* 220:262–271.
- Vuille, M., and M. Werner, 2005. Stable isotopes in precipitation recording South American summer monsoon and ENSO variability: Observations and model results. *Climate Dynamics* 25:401–413.
- Wagenführ, R., 2007. *Holzatlas*. Fachbuchverlag Leipzig im Carl Hanser Verlag, Germany.
- Watkinson, C. J., P. Gasson, G. O. Rees, and M. Boner, 2020. The development and use of isoscapes to determine the geographical origin of *Quercus* spp. in the United States. *Forests* 11:862. <https://doi.org/10.3390/f11080862>.
- West, J. B., G. J. Bowen, T. E. Cerling, and J. R. Ehleringer, 2006. Stable isotopes as one of nature's ecological recorders. *Trends in Ecology & Evolution* 21:408–414.
- West, J. B., G. J. Bowen, T. E. Dawson, and K. P. Tu, 2010. *Isoscapes: Understanding Movement, Pattern, and Process on Earth through Isotope Mapping*. Springer, Netherlands.
- Wickham, H., 2016. *ggplot2: Elegant Graphics for Data Analysis. R package version 2.2.1*. Springer-Verlag, New York.
- Wieloch, T., G. Helle, I. Heinrich, M. Voigt, and P. Schyma, 2011. A novel device for batch-wise isolation of α -cellulose from small-amount wholewood samples. *Dendrochronologia* 29: 115–117.
- Worbes, M., 1999. Annual growth rings, rainfall-dependent growth and long-term growth patterns of tropical trees from the Caparo Forest Reserve in Venezuela. *Journal of Ecology* 87:391–403.
- Zapater, M. A., E. M. Del Castillo, and T. D. Pennington, 2004. El genero *Cedrela* (Meliaceae) en la Argentina. *Darwiniana* 42:347–356.

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