

Ciencias Naturales

Artículo original

A Late Holocene Lecythidaceae wood record from Valle del Cauca, Colombia

Registro de madera de Lecythidaceae del Holoceno tardío en el Valle del Cauca, Colombia

 Diana Karen Pérez-Lara^{1,2,*},  Camila Martínez^{1,2},  Camila Pizano³,  Carlos Jaramillo²,  Víctor Vásquez¹

¹Área de Sistemas Naturales y Sostenibilidad, Universidad EAFIT, Medellín, Colombia

²Smithsonian Tropical Research Institute, Panamá City, Panamá

³Department of Biology, Lake Forest College, Illinois, USA

Abstract

A well-preserved fossil tree trunk was discovered in Valle del Cauca, Colombia. The radiocarbon date indicated that the tree lived during the Late Holocene, Meghalayan (4255 - 4083 cal BP). This finding is significant because the tree retains much of its original tissue. Based on wood anatomical comparisons, we suggest that this non-mineralized wood belongs to the family Lecythidaceae, a pantropical group of trees. This family includes around 12 recognized fossil genera based on fossil wood. The fossil probably belongs to the Bertholletia clade, which includes slow-growing, shade-tolerant species often found in tropical dry forests. This suggests that dry forests dominated the Valle del Cauca region before agricultural lands largely replaced them over the past decades.

Keywords: Meghalayan; Mummified; Dry forest; Subfossil; Non-mineralized.

Resumen

Se halló un tronco de árbol fósil bien conservado en el Valle del Cauca, Colombia. La fecha de radiocarbono indicó que el árbol vivió durante el Holoceno tardío, Meghalayo (4255 - 4083 cal BP). El descubrimiento es significativo porque el árbol conserva gran parte de su tejido original. Basándonos en comparaciones anatómicas de la madera, sugerimos que esta muestra no mineralizada pertenece a la familia Lecythidaceae, un grupo pantropical de árboles. Esta familia incluye alrededor de 12 géneros fósiles reconocidos a partir de maderas fósiles y el fósil encontrado probablemente pertenece al clado Bertholletia, el cual incluye especies de crecimiento lento, tolerantes a la sombra, comunes en los bosques secos tropicales, lo que sugiere que el Valle del Cauca estuvo dominado por este tipo de bosques antes de ser reemplazados por tierras agrícolas en las últimas décadas.

Palabras clave: Megalayense; Momificado; Bosque seco; Subfósil; No mineralizado.

Introduction

A well-preserved subfossil tree was discovered in Valle del Cauca, Colombia (**Figure 1**). The large tree trunk was found when drilling a well at a water treatment plant in Puerto Mallarino, Cali. The specimen exhibits remarkable preservation, with much of its original tissue still intact. This preservation is unusual, as most fossil samples are mineralized with silica or carbonates.

Subfossil wood can be found in surroundings that inhibit microbial decay and prevent exposure to oxygen and harsh chemicals. It can be mummified, retaining most of its original tissue; charcoalized due to combustion in an anaerobic environment, or coalified from intense heat and pressure during deep burial (**Mustoe, 2018**). Although most mummified

Citation: Pérez-Lara DK, *et al.* A Late Holocene Lecythidaceae wood record from Valle del Cauca, Colombia. Revista de la Academia Colombiana de Ciencias Exactas, Físicas y Naturales. 48(189):922-935, octubre-diciembre de 2024. doi: <https://doi.org/10.18257/raccefyn.3085>

Editor: Elizabeth Castañeda

*Corresponding autor:

Diana Karen Pérez-Lara;
dkperezl@eafit.edu.co

Received: October 9, 2024

Accepted: December 4, 2024

Published on line: December 11, 2024



This is an open access article distributed under the terms of the Creative Commons Attribution License.



Figure 1. Large tree trunk found at a water treatment plant in Puerto Mallarino, Valle del Cauca, Colombia

wood localities are found in Quaternary volcanic settings, there are also Neogene and Paleogene records from the Canadian Arctic, northwestern United States, Hungary, Italy, China, and Japan, among other sites (Mustoe, 2018).

These types of fossil fragments can be identified with similar methods to those used for modern wood preparation, where anatomical sections for microscopic observation are obtained with microtomes (Jagels *et al.*, 2005). We propose that this non-mineralized wood belongs to the Lecythidaceae family, a pantropical group of trees within the order Ericales (APG IV, 2016). This family is most diverse in the Neotropics (Mori *et al.*, 1990; Vargas & Dick, 2020) and consists of three subfamilies: Foetidioideae (Madagascar), Planchonioideae (Asia and Africa), and Lecythidoideae (Mori *et al.*, 2017), the latter being the largest with approximately 232 species (Vargas & Dick, 2020).

The Lecythidoideae crown clade dates back 46 million years ago, with the stem age at 62.7 million years (Vargas & Dick, 2020) suggesting that the family dispersed well after the breakup of Gondwana, and most of its major clades diversified during the Miocene (23-5.3 million years ago). Vargas & Dick (2020) proposed the Guyana floristic region as the ancestral range for many Lecythidoideae clades.

After Lecythidoideae was established in the Neotropics, it diversified into a wide range of species potentially shaped by Pleistocene refugia (Haffer, 1969; Thomas *et al.*, 2014); however, there is no comprehensive study on Lecythidoideae diversification. The multiple fossil records of Lecythidaceae have been primarily described from fossil pollen and wood and most of them are from India. The oldest record is from the Piauí Cretaceous in Brazil (**Table 1**).

Today, the Valle del Cauca region is almost entirely deforested and dominated by sugar cane crops. Only four Lecythidaceae species have been identified in its tropical dry forest: *Eschweilera caudiculata* R.Knuth 1939, *Gustavia speciosa* (Kunth) DC 1828, *Gustavia superba* (Kunth) O.Berg 1856, and *Lecythis minor* Jacq. 1763 (Pizano *et al.*, 2014; Silverstone-Sopkin, 2012).

Materials and methods

Geological setting

The fossil wood was collected at the Emcali Water Treatment Plant in Puerto Mallarino (**Figure 2**), near the Cauca River in Cali, department of Valle del Cauca, Colombia (3.4462 N; 76.4795 W at 1018 m.a.s.l.).

A three-meter-long tree trunk (**Figure 1**) was collected during the drilling of a 50-meter-deep water well at meter 17. In this area, Quaternary alluvial deposits are exposed (Gomez *et al.*, 2023), and probably the well crossed this particular one (**Figure 2**). The fossil was

Table 1. Lecythidaceae fossil species

Species	Vessels	Axial parenchyma	Rays	Crystals	Other	Region	Age	Reference
<i>Barringtonioxylon arcotense</i>	8–16/mm ²	Paratracheal and apotracheal diffuse	6–8/mm 1–8 seriate Homocellular / heterocellular	Absent	Tyloses and silice	South India	Paleogene	Awasthi, 1969
<i>Barringtonioxylon assamicum</i>	12–16/mm ²	Paratracheal parenchyma scanty, apotracheal diffuse	6–9/mm 1–7 seriate Homocellular / heterocellular	Absent	Tyloses and gums	Tipam sandstones Formation, India	Oligocene and Miocene	Prakash & Tripathi, 1970; Agarwal et al., 2000
<i>Barringtonioxylon eopterocarpum</i>	12/mm ²	Paratracheal parenchyma sparse, apotracheal abundant, occurring mostly as narrow, irregular, uniseriate lines	6–8/mm 1–8 seriate Homocellular / heterocellular	Absent	Tyloses	The Deccan Intertrappean beds of Mahurzari, India	Eocene	Prakash & Dayal, 1964
<i>Barringtonioxylon mandlaense</i>	---	Axial parenchyma diffuse vasicentric, aliform	4–12/mm 4–10 seriate Heterocellular	Absent	Sheath cells	The Deccan Intertrappean beds of Mandla District, India		Bande & Khatri, 1980
<i>Barringtonioxylon deccanense</i>	--	Diffuse-in-aggregates, scanty paratracheal and banded	4–12/mm >4-seriate Heterocellular	Absent	Axial canals, sheath cells, tyloses	India	Upper Cretaceous to Early Eocene	Shalom, 1960
<i>Careyoxylon pondicherriense</i>	8–15/mm ²	Paratracheal and apotracheal (1–6 cells wide)	9–14/mm 1–4 seriate Homocellular / heterocellular	Absent	Tyloses	South India	Paleogene	Awasthi, 1969
<i>Careyoxylon chindwinnense</i>	----	Diffuse-in-aggregates, banded in narrow bands up to three cells wide	4–12/mm 1–3 seriate Homocellular / heterocellular	Present	Tyloses	Myanmar	Middle Miocene	Gottwald, 1994
<i>Careyoxylon kuchilense</i>	6–14/mm ²	Paratracheal parenchyma scanty, apotracheal diffuse	7–15/mm 1–5 seriate Homocellular / heterocellular	—	Tyloses	Tipam sandstones Formation, India	Middle Miocene	Prakash & Tripathi, 1970
<i>Carinianoxylon brasiliensi</i>	7–14/mm ² 170µm	Reticulate	8–11/mm 1–3 seriate Homocellular / heterocellular	Present	Tyloses	Maranhão, Brazil Soliões Formation, Brazil	Paleogene Miocene	Selmeier, 2003 Kloster et al., 2017
<i>Lecythioxylon brasiliensi</i>	4–5/mm ²	In lines of 1–2 cells wide	7–12/mm 1–3 seriate Homocellular / Heterocellular	Present	—	Piauí, Brazil	Cretaceous	Milanez, 1935
<i>Lecythioxylon milanezzi</i>	4–5/mm ²	In lines of 1–2 cells wide	9–12/mm 1–3 seriate Homocellular / heterocellular	Present	Tyloses	Brazil	Neogene	Mussa, 1959
<i>Cariniana valverdei</i>	7/mm ²	Paratracheal scarce and apotracheal of 1–3 cells wide	5/mm 1–4 seriate Homocellular / heterocellular	Present	Tyloses	Piedra Chamaná Fossil Forest, Perú	Eocene	Woodcock et al., 2017

Species	Vessels	Axial parenchyma	Rays	Crystals	Other	Region	Age	Reference
<i>Lecythioxylon enviraense</i> , <i>cf. Cariniana</i>	4–15/mm ² 128µm	Reticulate >3 cells wide	7-12/mm 1-3 seriate Homocellular /heterocellular	Present	Tyloses	Solimões Formation, Brazil	Miocene	Kloster <i>et al.</i> , 2017
<i>cf. Echweilera</i>	--	Reticulate 1-2 cells wide	1-2 seriate Homocellular	Absent	Two types of vessel-ray pits are present	Pebas formation, Perú	Middle Miocene to Pliocene	Pons & de Franceschi, 2007
Lecythidaceae this work	6-12/mm ²	In broad continuous tangential bands	2-3 seriate Homocellular	Absent	Tyloses	Pebas formation, Perú	Middle Miocene to Pliocene	Pons & de Franceschi, 2007
		Paratracheal absent and apotracheal diffuse, and in narrow bands up to 5 cells wide	8-11/mm 1-2 seriate Homocellular /heterocellular	Present	Tyloses	Valle del Cauca, Colombia	Holocene	

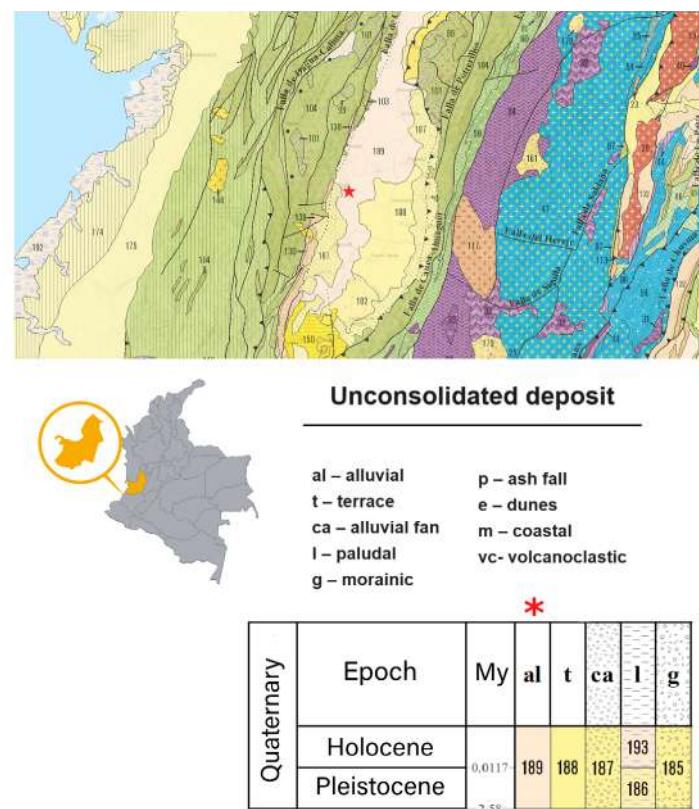


Figure 2. Sample collection site (red star) and stratigraphic column (red asterisk). Taken and modified from Gómez *et al.*, 2023.

sliced into 20 pieces with a benchtop band saw at Universidad ICESI (Cali, Colombia). Five of these pieces were sent to Universidad EAFIT (Medellín, Colombia), where they were sectioned for anatomical observation. Another fragment was sent to the Beta Analytic Radiocarbon Dating Laboratory (Miami, Florida, United States) for radiocarbon dating (sample ID 48530), and the remaining pieces remained at the Universidad ICESI Herbarium.

Sample processing

The anatomical features were studied from small pieces of wood found at the collection site and later placed in Ziploc bags to keep them moist until the time of preparation in the laboratory. Some of the well-preserved wood samples had to be dehydrated by immersion in 95% ethanol for 72 hours and then embedded in polyethyleneglycol (PEG) 2000 at 60°C for 24 hours. Then, we obtained 20-70 µm thick slices using a SLEE Mainz Rotary Microtome CUT 5062. The best-quality slices from each sample were immediately assembled using Eukitt to prevent the tissue from degrading.

We used the terminology from the IAWA Hardwood List (Wheeler, 1986; IAWA Committee, 1989) for the descriptions, and determined affinities by consulting the literature (de Zeeuw & Mori, 1987; Detienne & Jacquet, 1983; Diehl, 1935; Metcalfe & Chalk, 1950; Richter, 1982) and the multiple-entry key InsideWood from the Inside Wood Database (2004 onwards) (Wheeler, 2011). Photographs were taken with an Olympus BX53 light microscope and an SC100 digital camera with a 10.5 MP CMOS sensor.

We scanned the sample with a Scanning Electron Microscope to assess the type of non-mineralization (mummification, charcoalfication, or coalification) and the potential for stable isotope analyses (Mustoe, 2018: **Figure 3**). For the taxonomic classification, we followed APG IV (The Angiosperm Phylogeny Group, 2016).

Results

Radiometric dating

The radiocarbon dating of sample 48530 indicated a calibrated age of 4255 – 4083 cal BP (91.5%) (Beta Analytic 632015, Conventional Radiocarbon Age: 3790 ± 30 BP, percent modern carbon (pMC): 62.39 ± 0.23 pMC, $\delta_{13}\text{C}$ (IRMS): -25.3‰) (**Figure 4**).

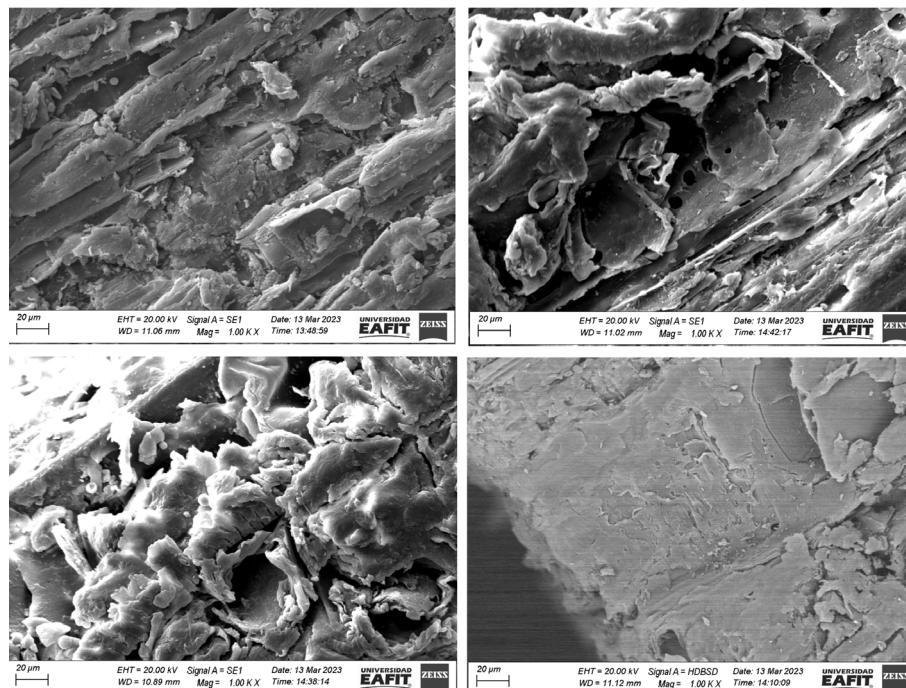


Figure 3. SEM photos. The wood's external structure remains largely preserved. The cell walls consist of an outer primary wall that encases a multi-layered secondary wall. The primary wall, which can be relatively thick, is primarily made up of cellulose and hemicellulose. In contrast, the thinner secondary wall is predominantly composed of cellulose and lignin. This secondary wall is typically divided into three distinct layers (lamellae), each varying in the arrangement of cellulose aggregates and the amount of lignin.

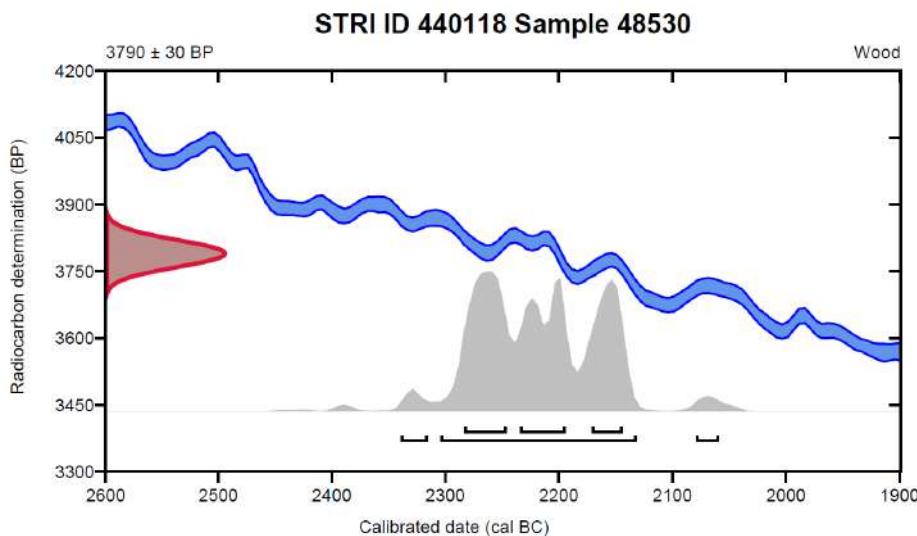


Figure 4. Calibration results using radiocarbon ages

Systematic palaeontology

Order – Ericales

Family – Lecythidaceae

Subfamily — Lecythidoideae Beilschmied

Clade — Bertholletia

Genus — *Eschweilera/Lecythis*

Age –calibrated age of 4255 – 4083 cal BP (radiocarbon dating analysis obtained for the fossil fragment)

Description in IAWA feature numbers: 5p 13p 22p 26p 32p 56p 65p 66p 70p 87p 89p 97p 106p 115p with 1 allowable mismatch

Description – Growth rings indistinct. Diffuse porous wood (**Figure 5 a-c**). Solitary vessels (36%), radial multiples two to three (up to 6) (**Figure 5 a-c**). Vessel elements round to oval in outline (**Figure 5 b-c**). Mean tangential vessel diameter 102 (70-200) µm; 8 (6–12) vessels per square millimeter; vessel walls in transverse sections of 12 (6-16) µm thick; simple perforation plates; intervessel pitting alternate 7-10 µm (**Figure 5 d**); vessel-ray parenchyma pitting with reduced borders, horizontally elongate, and similar to intervacular pits (**Figure 5 h**). Mean vessel element length 200-400 µm. Bubble-like tyloses common (**Figure 5 b-c**).

Uniseriate rays averaging 12 (7–20) cells, 535 (407-1188) µm high, 22 (19-23) µm wide (**Figure 5 e-f**). Some biserrate rays averaging 12 (8–21) cells and 904 (710-1210) µm high, 28 (27-29) µm wide (**Figure 5 e-f**). Rays 10 (8–11) per millimeter. Heterogeneous with procumbent body cells and usually one marginal row (**Figure 5 g**).

Fibers septate (**Figure 5**) and non-septate, thin to thick walls, pitting not observed. Axial parenchyma apotracheal diffuse, and axial parenchyma in narrow bands up to five cells wide (**Figure 5 a-c**).

Remarks — The fossil wood studied exhibits a combination of features indicating an affinity with Lecythidaceae and Malvaceae. However, Malvaceae members generally have storied rays and scarce axial parenchyma (**Wheeler et al.**, 1994). This fossil wood shows characters consistent with Lecythidaceae, such as parenchyma in apotracheal bands, simple perforation plates, alternate intervessel pitting, and two types of vessel-ray pitting. Additionally, rays are mostly 2–3 cells wide and distinctly heterogeneous to homogeneous, which aligns with the characteristics of Lecythidaceae (**Diehl**, 1935; **Lens et al.**, 2007;

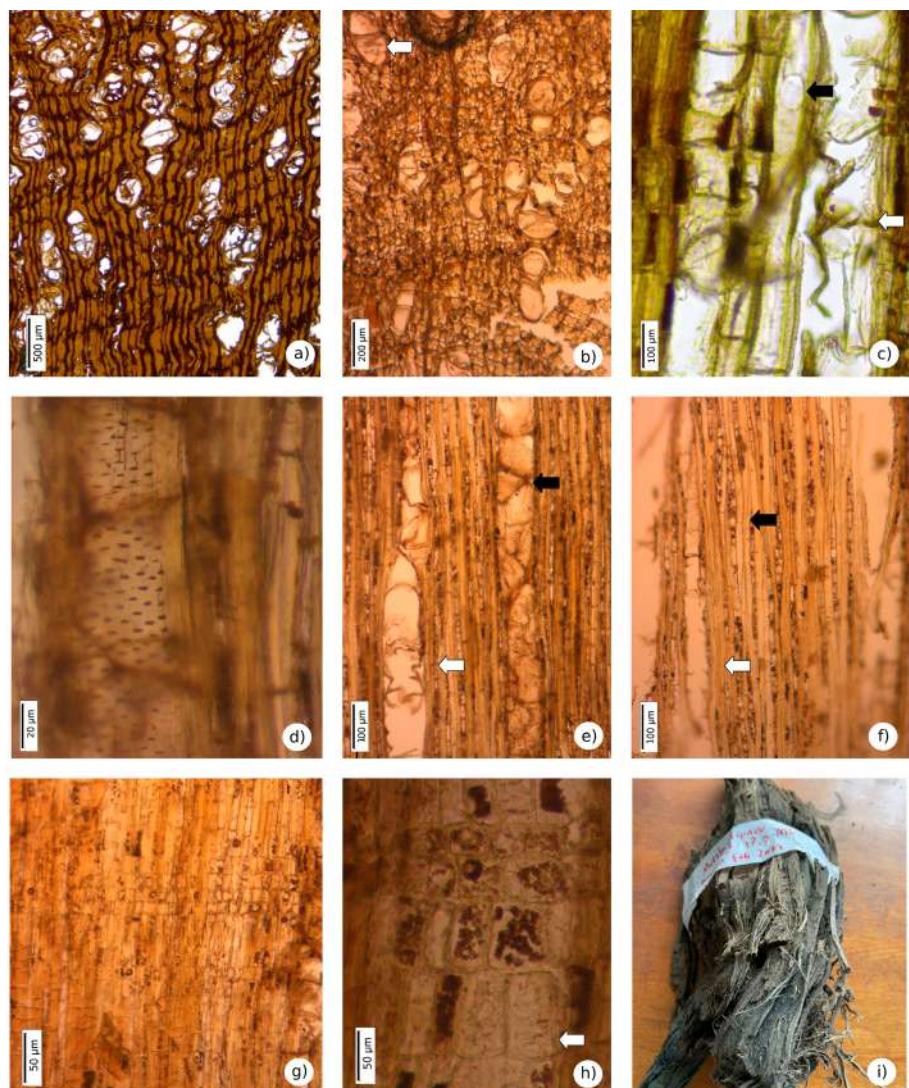


Figure 5. Wood anatomical features of the mummified wood. **a.** Transverse section (TS). Diffuse porous wood. **b.** TS. Vessels in radial multiples and nearly absent parenchyma. **c.** Axial parenchyma apotracheal diffuse and axial parenchyma in narrow bands up to five cells wide. **d.** Longitudinal tangential section (LTS). Vessel elements, simple perforation plates, and alternate intervessel pitting. **e.** LTS. Tyloses and uniseriate rays and biserrate rays. **f.** LTS. Septate and non-septate fibers and uniseriate rays. **g.** LRS. Heterogeneous with procumbent body cells and usually one marginal row. **h.** Vessel-ray parenchyma pitting with reduced borders, horizontally elongated and similar to intervacular pits. **i.** mummified wood.

Metcalfe & Chalk, 1950; Silva et al., 2022). Rays < 1 mm high lacking crystals and the presence of two types of vessel-ray pits generally distinguish the Lecythidoideae subfamily (Lens et al., 2007). The solitary vessels in small multiple tyloses, the parenchyma in continuous bands 1–2 cells wide, and the abundant less than 1 mm uniseriate rays with 1 or 2 marginal rows of square cells support its inclusion in *Eschweilera* and *Lecythis* (Metcalfe & Chalk, 1959) (**Tabla 2**).

Comparisons with Lecythidaceae fossil woods

Approximately 12 fossil woods have been described (Gregory et al., 2009; Inside Wood Database, 2004-onwards) (**Table 1**). The following characteristics differentiate them

Table 2. Some modern Lecythidaceae species

Species	Vessels	Axial parenchyma	Rays	Crystals	Other
<i>Gustavia speciosa</i> Kunth	1–30/mm ²	Diffuse-in aggregates to uniseriately banded	1-11/mm 2-8 seriate Heterocellular	Present	_____
<i>Gustavia augusta</i> L.	20-40/mm	Diffuse, diffuse-in aggregates, scanty, narrow bands	6-10/mm 1-9 seriate Heterocellular	In chambered axial parenchyma cells	Tyloses
<i>Gustavia gigantophylla</i> Sandwith	5-20/mm	Narrow bands	4-12/mm 4-10 seriate Heterocellular	In chambered axial parenchyma cells	Tyloses
<i>Gustavia hexapétala</i> (Aubl.) Sm.	20-100/mm	Diffuse-in aggregates	4-12/mm 1-3 seriate Heterocellular	In chambered axial parenchyma cells and in rays	_____
<i>Gustavia nana</i> Pitt.	20-40/mm	Narrow bands	4-12/mm >10 seriate Heterocellular	In chambered axial parenchyma cells	
<i>Lecythis ampla</i>	8-15/mm ²	Paratracheal and apotracheal (1-6 cells wide)	9-14/mm 1-4 seriate Homocellular/heterocellular	Absent	Tyloses
<i>Lecythis chartacea</i> O.Berg	5-20/mm	Bands (> 3 cells)	4-12/mm 1-7 Homocellular to heterocellular	In chambered axial parenchyma cells	Tyloses
<i>Lecythis confertiflora</i> (A.C.Sm.) S.A.Mori	<5-20/mm	Scanty paratracheal, narrow bands, reticulate, marginal bands	4-12/mm 1-3 seriate Homocellular to heterocellular	In chambered axial parenchyma cells	Tyloses and silica bodies in rays
<i>Lecythis idatimon</i> Aubl.	<5-20/mm	Scanty paratracheal, narrow bands, and reticulate	4-12/mm 1-3 Homocellular	_____	Tyloses
<i>Lecythis lurida</i> (Miers) S.A.Mori	<5-20/mm	Diffuse, diffuse-in aggregates	4-12/mm 1-6 Heterocellular	In chambered axial parenchyma	Tyloses
<i>Lecythis pisonis</i> Cambess.	<5-20/mm	Narrow bands, reticulate, marginal bands	4-12/mm 1-3 Homocellular to heterocellular	In chambered axial parenchyma	Tyloses and silica bodies in rays
<i>Lecythis zabucajo</i> Aubl.	<5-20/mm	Scanty, narrow bands, reticulate and marginal bands	4-12 1-3 Homocellular to heterocellular	In chambered axial parenchyma	Tyloses and silica bodies in rays
<i>Eschweilera sclerophylla</i> Cuatrec	7-14/mm ² 170µm	Reticulate	8-11/mm 1-3 seriate Homocellular/heterocellular	Present	Tyloses
<i>Eschweilera alata</i> A.C.Sm.	<5/mm	Scanty, narrow bands, reticulate and in marginal bands	10-14/mm 1-3 seriate Homocellular to heterocellular	_____	Tyloses and silica bodies in rays
<i>Eschweilera tenax</i> (Moritz ex O.Berg) Miers	5-20/mm	Bands and reticulate	4-12/mm 1-3 seriate Homocellular	In chambered axial parenchyma cells	Tyloses
<i>Eschweilera albiflora</i>	5-20/mm	Bands, reticulate	4-12/mm 1-4 seriate Homocellular to weakly heterocellular	In chambered axial parenchyma cells	Tyloses
<i>Eschweilera congestiflora</i>	5-20/mm	Bands	4-12/mm 1-4 seriate Heterocellular	_____	Tyloses
Lecythidaceae this work	6-12/mm ²	Apotracheal diffuse and paratracheal in narrow bands	8-11/mm 1-2 seriate Homocellular/heterocellular	In axial parenchyma	Tyloses

from the fossil we describe here: *Barringtonioxylon arcotense* Awasthi, 1969 from the Indian Neogene exhibits parenchyma aliform to confluent and wider rays (up to 8 cells) (**Awasthi**, 1969). *Barringtonioxylon eopterocarpum* from the Eocene of the Deccan Intertrappean Series, India, differs from our wood sample in that it has ring-porous wood and lacks radial canals (**Prakash & Dayal**, 1964). *Barringtonioxylon mandlaense* from the Deccan Intertrappean Series, India, has vasicentric parenchyma and fine to broad rays of two distinct types (**Bande & Khatri**, 1980). *Careyoxylon pondicherriense* reported in Paleogene sediments from southern India differs in having wider rays (up to 4 cells) and sheath cells (**Awasthi**, 1969). *Careyoxylon chindwinnense* from the middle Myanmar Miocene differs in having prismatic crystals in rays and thick-walled vessels (**Gottwald**, 1994; *Inside Wood Database*, 2004-onwards). *Careyoxylon kuchilense* described from the Tipam sandstones Formation Middle Miocene in India differs in the presence of scanty paratracheal parenchyma (**Prakash & Tripathi**, 1970). In the New World, *Carinianoxylon brasiliensi* from Brazil differs in having reticulate parenchyma (**Kloster et al.**, 2017; **Selmeier**, 2003). *Lecythioxylon brasiliensi* and *Lecythioxylon milanezzi* described from the Neogene have lower vessel density and wider rays (**Milanez**, 1935; **Mussa**, 1959). *Lecythioxylon enviraense*, from the Acre Basin, Brazil, differs in having reticulate axial parenchyma more than three cells wide and the presence of crystals in ray cells (**Kloster et al.**, 2017). Finally, from the Eocene of Perú, *Cariniana valverdei* has lower vessel density and scarce paratracheal parenchyma (**Woodcock et al.**, 2017) (**Table 1**).

Scanning electron microscopy

Photographs were taken to examine the anatomy and preservation of the samples (**Figure 3**). The tangential section is shown in greater detail, revealing key anatomical features such as fibers, parenchyma, vessels, and intervessel pits (**Figure 5**).

Remarks — **Mustoe** (2018) identified three types of non-mineralized fossilized wood: mummified, charcoalfied, and coalified. Mummified wood retains its original tissues with minimal chemical alteration, though they may exhibit distortion or desiccation. Charcoalfied wood forms when wood is burned in an oxygen-deprived environment, reducing the organic material mainly to pure carbon. Coalified wood, in contrast, results from prolonged exposure to heat and pressure during deep burial, transforming the original organic material into a mixture of pure carbon and hydrocarbons.

Discussion

Subfossil wood has an economic importance in the wood industry, especially in Europe, where it is used in furniture (**Beldean & Timar**, 2021). In Colombia, information about subfossil wood is scarce — only two studies describe five subfossil woods identified as *Terminalioxylon gumminae*, *Andesanthus risaraldense*, *Anacardium quindiuense*, *Chrysochlamys colombiana*, and *Goupioxylon stutzeri* from the Pleistocene of the Central Cordillera (**Ayala-Usma**, 2014; **Ayala-Usma et al.**, 2024).

The successful preparation, description, and dating of this sample confirms the high potential that non-mineralized fossil wood has for paleobotanical investigation. The processes described here can be replicated for future findings of this type of fossil. Our record confirms the presence of the Bertholletia clade (*Lecythis/Eschweilera*) in Valle del Cauca at least since the Late Holocene (Meghalayan).

This wood is an exemplar of mummified wood because it retains its original tissues with only minimal degradation of cellular components (**Figures 1, 3, 5i**) (**Mustoe**, 2018). The key conditions for mummification involve preventing microbial and chemical decay, typically in environments with deeply submerged wood, burial in impermeable sediments, arid regions, or cold climates. Also, this subfossil wood is frequently found in Quaternary deposits, where glacial, fluvial, and volcaniclastic lahar sediments create deep layers of fine material that aid in preserving organic remains (**Mustoe**, 2018).

Cali, where the fossil was found, is the capital of the department of Valle del Cauca and the third largest city in Colombia, with a population of approximately 2.28 million (**DANE**, 2018) and an average elevation of 1018 m. The surrounding area is highly urbanized, the Cauca River valley has been largely transformed, and only 1.4% of primary forest, mostly tropical dry forest, remains (**Pizano et al.**, 2014).

The presence of the Lecythidaceae family in Colombia had been already reported. **Berrio et al.** (2002) described *Couroupita santanderiensis* fossil leaves from the Miocene of Santander and **Huertas** (1969) described a *Lecythisdopyon girardotanum* fruit from Cundinamarca.

Lecythis and *Eschweilera* are non-monophyletic genera that share the greatest number of characters with our fossil and belong to the Lecythidoideae subfamily, which is dominant in the Amazonian forests and includes many emergent tree species (**Vargas & Dick**, 2020). Both genera vary from small to very large trees found throughout the Neotropics below 1500 m (**GBIF**, 2024) in México, Central America, Colombia, Venezuela, Amazonia, and Eastern and Central Brazil (**Huang et al.**, 2015; **Mori et al.**, 2017, 2007). The genera originated in the biogeographic regions of the Guiana, Amazonia, Cerrado-Caatinga, Mata Atlántica, and Transandean areas (**Vargas & Dick**, 2020). It is possible then that these species were typical of the dry forest-wetland transition that characterized this vast region before natural ecosystems were almost completely replaced by agricultural lands (**Ramos-Pérez & Silverstone-Sopkin**, 2018; **Silverstone-Sopkin**, 2012).

Late Holocene climate

The Valle del Cauca vegetation was dominated by tropical dry forests before the Industrial Revolution. The climate of the late Holocene, around 4,200 years BP, was a period characterized by increased aridity in mid- and low-latitudes (Meghalayan) (**Berkelhammer et al.**, 2012; **Walker et al.**, 2018, 2012). Many records show a rapid onset of aridification in the Mediterranean region, parts of Asia, North America, northeastern Brazil, and Africa (**Bini et al.**, 2019; **Booth et al.**, 2005; **Cheng et al.**, 2015; **De Oliveira et al.**, 1999; **Dixit et al.**, 2014; **Kaniewski et al.**, 2017; **Thompson et al.**, 2002; **Toth & Aronson**, 2019; **Utida et al.**, 2019, 2020). In Colombia, **Berrio et al.** (2002) described two sediment cores from the Valle del Cauca region with pollen, charcoal, and radiocarbon data. The Quilichao-1 core covers from 13,150 to 7,720 14C years BP and, after a gap, from 2,880 14C years BP to the present. The La Teta-2 core provides a continuous record from 8,700 14C years BP to the present. Maximum dryness was reached around 7,500 and 4,300 14C years BP, close to the 4,200 events. Additionally, pollen grains from Zone TET-1 (8850–7560 14C yr BP) show the dominance of dry forest taxa (*Crotalaria*, *Croton*, *Tabebuia*, *Alchornea*), while Lecythidaceae represents 8% of the pollen grains. Although the Valle del Cauca records for the late Holocene indicate an increase in aridity, the changes in precipitation were not large enough to modify its biome dominance; the valley was dominated by dry forests and wetlands/flooded forests during the Late Holocene and in modern times before the Industrial Revolution.

Conclusions

The discovery of a well-preserved subfossil tree in Valle del Cauca, Colombia, provides a rare and valuable insight into the ancient tropical dry forests of the region. The fossil wood exhibits anatomical features strongly indicative of the Lecythidaceae family, particularly aligning with the genera *Lecythis* and *Eschweilera*. These findings suggest that Lecythidaceae, a family now dominant in the Amazon forests, once flourished in the Valle del Cauca region, contributing to the region's biodiversity and ecological complexity. They further underscore that the region probably lost a considerable part of its biodiversity centuries ago, when natural forests were replaced by agricultural lands (**Silverstone-Sopkin**, 2012). The identification of Lecythidaceae in the fossil record highlights the long-term ecological stability and adaptability of this family. Furthermore, the preservation of this fossil wood suggests that specific environmental conditions, such as rapid burial

and low oxygen levels, played a crucial role in the fossilization of wood. The climatic history of the Valle del Cauca region is a powerful reminder of the capacity of natural systems to adapt and evolve in the face of change, but also of their vulnerability. Learning from these records can help us anticipate future challenges and more effectively adapt our conservation and climate management strategies.

Acknowledgments

We want to thank EMCALI-ECIE-ESP for the permission to collect and study the fossil. Luis Darío Gómez (CINARA-Universidad del Valle) and Juan Pablo Gutiérrez and José Ospinal (MCALI-ECIE-ESP) for making this study possible; Wilmar Oswaldo Díaz-Vasco for helping with the field collection and the processing and proper storage of the fossil at the Universidad Icesi Herbarium; Carlos D. Cárdenas and Juan David Quintero for their assistance in the processing of the fossil, and Juan Camilo Arango Gálvez for their assistance in taking the SEM photographs at Universidad EAFIT.

Conflicts of interest

The authors declare that they have no competing financial interests or personal relationships that could have influenced the work reported in this paper.

References

- Agarwal, A., Ambwani, K., Saha, S., and Kar, R.K.** (2000). Fossil wood of Barringtonia (Lecythidaceae) from Ramgarh, Chittagong Hill Tract, Bangladesh. *Phytomorphology*, 50, 333-336.
- Awasthi, N.** (1969). On the occurrence of two new fossil woods belonging to the family Lecythidaceae in the Tertiary rocks of South India. *Journal of Palaeosciences*, 18(1-3), Article 1-3. <https://doi.org/10.54991/jop.1969.820>
- Ayala-Usma, D. A.** (2014). *Stable isotopes in subfossil woods-first insights into an ultra-high resolution paleoclimatic record of an Andean Forest during the Late Pleistocene*. <https://repositorio.uniandes.edu.co/bitstreams/adcb7ee4-5a14-4b57-a138-7cea0817ec9b/download>
- Ayala-Usma, D. A., Lozano-Gutiérrez, R., Orejuela, C., Pérez-Ángel, L. C., Montes, C., González-Arango, C.** (2024). Exceptionally preserved subfossil woods from late Pleistocene volcanic deposits from the Northern Andes of Colombia. *Review of Palaeobotany and Palynology*, 324, 105090. <https://doi.org/10.1016/j.revpalbo.2024.105090>
- Bande, M. B. & Khatri, S. K.** (1980). Some more fossil woods from the Deccan intertrappean beds of Mandla District, Madhya Pradesh, India. *Palaeontographica Abteilung B Band*, 173(4-6), 147-165.
- Beldean, E. & Timar, M. C.** (2021). A New Opportunity for Research in Romania – Subfossil Wood. *Bulletin of the Transilvania University of Brasov. Series II: Forestry • Wood Industry • Agricultural Food Engineering*, 77-88. <https://doi.org/10.31926/but.fwiafe.2021.14.63.1.7>
- Berkelhammer, M., Sinha, A., Stott, L., Cheng, H., Pausata, F. s. r., Yoshimura, K.** (2012). An Abrupt Shift in the Indian Monsoon 4000 Years Ago. *Climates, Landscapes, and Civilizations*. American Geophysical Union (AGU). <https://doi.org/10.1029/2012GM001207>
- Berrío, J. C., Hooghiemstra, H., Marchant, R., Rangel, O.** (2002). Late-glacial and Holocene history of the dry forest area in the south Colombian Cauca Valley. *Journal of Quaternary Science*, 17(7), 667-682. <https://doi.org/10.1002/jqs.701>
- Bini, M., Zanchetta, G., Perşoiu, A., Cartier, R., Català, A., Cacho, I., Dean, J. R., Di Rita, F., Drysdale, R. N., Finnè, M., Isola, I., Jalali, B., Lirer, F., Magri, D., Masi, A., Marks, L., Mercuri, A. M., Peyron, O., Sadori, L., ... Brisset, E.** (2019). The 4.2 ka BP Event in the Mediterranean region: An overview. *Climate of the Past*, 15(2), 555-577. <https://doi.org/10.5194/cp-15-555-2019>
- Booth, R. K., Jackson, S. T., Forman, S. L., Kutzbach, J. E., Bettis, E. A., Kreigs, J., Wright, D. K.** (2005). A severe centennial-scale drought in midcontinental North America 4200 years ago and apparent global linkages. *The Holocene*, 15(3), 321-328. <https://doi.org/10.1191/0959683605hl825ft>

- Cheng, H., Sinha, A., Verheyden, S., Nader, F. H., Li, X. L., Zhang, P. Z., Yin, J. J., Yi, L., Peng, Y. B., Rao, Z. G., Ning, Y. F., Edwards, R. L.** (2015). The climate variability in northern Levant over the past 20,000 years. *Geophysical Research Letters*, 42(20), 8641-8650. <https://doi.org/10.1002/2015GL065397>
- DANE** (2018). *¿Cuántos somos?* <https://www.dane.gov.co/index.php/estadisticas-por-tema/demografia-y-poblacion/censo-nacional-de-poblacion-y-vivenda-2018/cuantos-somos>
- De Oliveira, P. E., Barreto, A. M. F., Suguio, K.** (1999). Late Pleistocene/Holocene climatic and vegetational history of the Brazilian caatinga: The fossil dunes of the middle São Francisco River. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 152(3), 319-337. [https://doi.org/10.1016/S0031-0182\(99\)00061-9](https://doi.org/10.1016/S0031-0182(99)00061-9)
- de Zeeuw, C. H. & Mori, S. A.** (1987). Wood anatomy. The Lecythidaceae of a lowland Neotropical forest: La Fumée Mountain, French Guiana. *Memoirs of the New York Botanical Garden*, 44, 100-112.
- Detienne, P. & Jacquet, P.** (1983). *Identification atlas for wood of the Amazon River and neighboring areas*. Centre technique forestier tropical. https://scholar.google.com/?title=Identification+atlas+for+wood+of+the+Amazon+River+and+neighboring+areas&author=Detienne%2C+Pierre.&publication_year=1983
- Diehl, G. A.** (1935). *A study of the Lecythidaceae*. Yale University School of Forestry.
- Dixit, Y., Hodell, D. A., Petrie, C. A.** (2014). Abrupt weakening of the summer monsoon in northwest India ~4100 yr ago. *Geology*, 42(4), 339-342. <https://doi.org/10.1130/G35236.1>
- GBIF Backbone Taxonomy** (2024). <https://doi.org/10.15468/39omei>
- Gómez, J., Montes, N. E., Marín, E.** (2023). *Mapa Geológico de Colombia 2023*. <https://sgcolumbiano.maps.arcgis.com/apps/mapviewer/index.html?webmap=6faed511e7c542de85ae4d6a1668953c>
- Gottwald, H. P.** (1994). *Tertiäre Kieselhölzer aus dem Chindwinn-Bassin im nordwestlichen Myanmar (Birma)*. Forschungen aus den Naturwissenschaften.
- Gregory, M., Poole, I., Wheeler, E. A.** (2009). *Fossil Dicot Wood Names: An Annotated List with Full Bibliography*. Published for the International Association of Wood Anatomists at the Nationaal Herbarium Nederland.
- Haffer, J.** (1969). Speciation in Amazonian Forest Birds. *Science*, 165(3889), 131-137. <https://doi.org/10.1126/science.165.3889.131>
- Huang, Y.-Y., Mori, S. A., Kelly, L. M.** (2015). Toward a phylogenetic-based Generic Classification of Neotropical Lecythidaceae—I. Status of Bertholletia, Corythophora, Eschweilera and Lecythis. *Phytotaxa*, 203(2), 85. <https://doi.org/10.11646/phytotaxa.203.2.1>
- Huertas G. G.** (1969). Un nuevo género y especie fósiles de las Lecitidaceas. *Caldasia*, 10(48), 365-369.
- Jagels, R., Visscher, G.E., Wheeler, E.A.** (2005). An Eocene High Arctic Angiosperm Wood. *IAWA Journal*, 26(3), 387-382. <https://doi.org/10.1163/22941932-02603009>
- Jansen, S., Kitin, P., De Pauw, H., Idris, M., Beeckman, H., Smets, E.** (1998). Preparation of Wood Specimens for Transmitted Light Microscopy and Scanning Electron Microscopy. *Belgian Journal of Botany*, 131(1), 41-49.
- Kaniewski, D., Marriner, N., Ilan, D., Morhange, C., Thareani, Y., Van Campo, E.** (2017). Climate change and water management in the biblical city of Dan. *Science Advances*, 3(11), e1700954. <https://doi.org/10.1126/sciadv.1700954>
- Kloster, A., Gnaedinger, S., Adami-Rodrigues, K.** (2017). Nuevos registros de maderas del Mioceno de la Formación Solimões, Cuenca de Acre, Amazonia, Brasil. *Revista Brasileira de Paleontología*, 20(3), 383-404. <https://doi.org/10.4072/rbp.2017.3.09>
- Lens, F., Baas, P., Jansen, S., Smets, E.** (2007). A search for phylogenetically informative wood characters within Lecythidaceae s.l. *American Journal of Botany*, 94(4), 483-502. <https://doi.org/10.3732/ajb.94.4.483>
- Metcalfe, C. R. & Chalk, L.** (1950). *Anatomy of the dicotyledons, Vols. 1 & 2*. <https://www.cabidigitallibrary.org/doi/full/10.5555/19511603379>
- Milanez, F. R.** (1935). Estudo De Um Dicotyledoneo Fossil Do Cretaceo. *Rodriguésia*, 1(2), 83-89.
- Mori, S. A., Prance, G. T., De Zeeuw, C., Prance, G. T.** (1990). Zygomorphic-flowered New World genera (*Couroupita*, *Corythophora*, *Bertholletia*, *Couratari*, *Eschweilera*, & *Lecythis*). *Flora Neotropica*, 21, 1-333.
- Mori, S. A., Tsou, C.-H., Wu, C.-C., Cronholm, B., Anderberg, A. A.** (2007). Evolution of Lecythidaceae with an emphasis on the circumscription of neotropical genera: Information from combined ndhF and trnL-F sequence data. *American Journal of Botany*, 94(3), 289-301. <https://doi.org/10.3732/ajb.94.3.289>

- Mori, S., Kiernan, E., Smith, N., Kelly, L., Prance, G. T.** (2017). Observations on the phytogeography of the Lecythidaceae clade (Brazil nut family). *Phytoneuron*, 30, 1-85.
- Mussa, D.** (1959). *Contribuição à paleoanatomia vegetal: II madeiras fósseis do Território do Acre (Alto Juruá) Brasil*. Serviço Gráfico do Instituto Brasileiro de Geografia e Estatística.
- Mustoe, G. E.** (2018). Non-Mineralized Fossil Wood. *Geosciences*, 8(6), Article 6. <https://doi.org/10.3390/geosciences8060223>
- Pizano, C., Cabrera, M., García, H.** (2014). *Bosque seco tropical en Colombia; generalidades y contexto*. Bogotá, D.C. Instituto de Investigación de Recursos Biológicos Alexander von Humboldt (IAvH). <https://dare.uva.nl/search?identifier=f6d3da1b-126f-41a1-9040-1969c4c294ee>
- Pons, D., Franceschi, D.** (2007). Neogene woods from western Peruvian Amazon and paleoenvironmental interpretation. *Bulletin of Geoscience*, 82, 343-354. doi:10.3140/bull.geosci.2007.04.343
- Prakash, U. & Dayal, R.** (1964). Barringtonioxylon copterocarpum sp. Nov., A fossil wood of lecythidaceae from the Deccan Intertrappean beds of Mahurzari. *Journal of Palaeosciences*, 13(1-3), Article (1-3). <https://doi.org/10.54991/jop.1964.675>
- Prakash, U. & Tripathi, P. P.** (1970). Fossil woods of Careya and Barringtonia from the Tertiary of Assam. *Journal of Palaeosciences*, 19(1-3), 25-29. <https://doi.org/10.54991/jop.1970.863>
- Ramos-Pérez, J. E. & Silverstone-Sopkin, P. A.** (2018). *Catálogo de la Flora Relictual del Valle Geográfico del Río Cauca [Catalog of the Relictual Flora of the Geographical Valley of the Cauca River]*. <https://www.nhbs.com/catalogo-de-la-flora-relictual-del-valle-geografico-del-rio-cauca-catalog-of-the-relictual-flora-of-the-geographical-valley-of-the-cauca-river>
- Richter, H. G.** (1982). The Wood Structure of *Couratari* Aubl. and *Couroupita* Aubl. (Lecythidaceae). *IAWA Journal*, 3(1), 45-55. <https://doi.org/10.1163/22941932-90000746>
- Selmeier, A.** (2003). First record of a Lecythidaceae wood (*Carinianoxylon brasiliense* gen. Et sp. Nov.) from the Tertiary of the New World (Brazil, Rio Paranaiba). *Zitteliana*, 43, 171-178. <http://epub.ub.uni-muenchen.de/12128/>
- Shallom, L.J.** (1960). Fossil dicotyledonous wood of Lecythidaceae from the Deccan Intertrappean beds of Mahurzari. *Indianbot Soc.* 39(2), 198-203.
- Silva, M. dos S., Silva, L. B. da, Santos, N. C. dos, Carvalho, E. R. de, Silva, C. R. A. da, Silva, C. C. S., Leite, K. R. B.** (2022). *Madeiras da Bahia: Anatomia do lenho de espécies nativas da Mata Atlântica*. EDUFBA. <https://repositorio.ufba.br/handle/ri/34749>
- Silverstone-Sopkin, P. A.** (2012). *Los muertos vivientes: La historia natural de cuatro lirios amazónicos del suroccidente de Colombia (Eucharis y Plagiolirion, Amaryllidaceae)* (1.^a ed.). Universidad del Valle. <https://www.jstor.org/stable/j.ctv1228j4j>
- The Angiosperm Phylogeny Group.** (2016). An update of the Angiosperm Phylogeny Group classification for the orders and families of flowering plants: APG IV. *Botanical Journal of the Linnean Society*, 181(1), 1-20. <https://doi.org/10.1111/boj.12385>
- Thomas, E., Alcazar Caicedo, C., Loo, J., Kindt, R.** (2014). *The distribution of the Brazil nut (*Bertholletia excelsa*) through time: From range contraction in glacial refugia to anthropogenic climate change*. Boletín del Museo Paraense Emilio Goeldi. Ciencias Naturales. <https://hdl.handle.net/10568/6607>
- Thompson, L. G., Mosley-Thompson, E., Davis, M. E., Henderson, K. A., Brecher, H. H., Zagorodnov, V. S., Mashiotta, T. A., Lin, P.-N., Mikhalenko, V. N., Hardy, D. R., Beer, J.** (2002). Kilimanjaro Ice Core Records: Evidence of Holocene Climate Change in Tropical Africa. *Science*, 298(5593), 589-593. <https://doi.org/10.1126/science.1073198>
- Toth, L. T. & Aronson, R. B.** (2019). The 4.2 ka event, ENSO, and coral reef development. *Climate of the Past*, 15(1), 105-119. <https://doi.org/10.5194/cp-15-105-2019>
- Utida, G., Cruz, F. W., Etourneau, J., Bouloubassi, I., Scheuß, E., Vuille, M., Novello, V. F., Prado, L. F., Sifeddine, A., Klein, V., Zular, A., Viana, J. C. C., Turcq, B.** (2019). Tropical South Atlantic influence on Northeastern Brazil precipitation and ITCZ displacement during the past 2300 years. *Scientific Reports*, 9(1), 1698. <https://doi.org/10.1038/s41598-018-38003-6>
- Utida, G., Cruz, F. W., Santos, R. V., Sawakuchi, A. O., Wang, H., Pessenda, L. C. R., Novello, V. F., Vuille, M., Strauss, A. M., Borella, A. C., Strikis, N. M., Guedes, C. C. F., Dias De Andrade, F. R., Zhang, H., Cheng, H., Edwards, R. L.** (2020). Climate changes in Northeastern Brazil from deglacial to Meghalayan periods and related environmental impacts. *Quaternary Science Reviews*, 250, 106655. <https://doi.org/10.1016/j.quascirev.2020.106655>

- Vargas, O. M. & Dick, C. W.** (2020). Diversification History of Neotropical Lecythidaceae, an Ecologically Dominant Tree Family of Amazon Rain Forest. En V. Rull & A. C. Carnaval (Eds.), *Neotropical Diversification: Patterns and Processes* (pp. 791-809). Springer International Publishing. https://doi.org/10.1007/978-3-030-31167-4_29
- Walker, M., Head, M. J., Berkelhammer, M., Björck, S., Cheng, H., Cwynar, L., Fisher, D., Gkinis, V., Long, A., Lowe, J., Newnham, R., Rasmussen, S. O., Weiss, H.** (2018). Formal ratification of the subdivision of the Holocene Series/Epoch (Quaternary System/Period): Two new Global Boundary Stratotype Sections and Points (GSSPs) and three new stages/subseries. *Episodes Journal of International Geoscience*, 41(4), 213-223. <https://doi.org/10.18814/epiugs/2018/018016>
- Walker, M. J. C., Berkelhammer, M., Björck, S., Cwynar, L. C., Fisher, D. A., Long, A. J., Lowe, J. J., Newnham, R. M., Rasmussen, S. O., Weiss, H.** (2012). Formal subdivision of the Holocene Series/Epoch: A Discussion Paper by a Working Group of INTIMATE (Integration of ice-core, marine and terrestrial records) and the Subcommission on Quaternary Stratigraphy (International Commission on Stratigraphy). *Journal of Quaternary Science*, 27(7), 649-659. <https://doi.org/10.1002/jqs.2565>
- Wheeler, E. A.** (1986). Vessels Per Square Millimetre or Vessel Groups Per Square Millimetre? *IAWA Journal*, 7(1), 73-74. <https://doi.org/10.1163/22941932-90000444>
- Wheeler, E. A.** (2011). Inside Wood – A Web resource for hardwood anatomy. *IAWA Journal*, 32(2), 199-211. <https://doi.org/10.1163/22941932-90000051>
- Wheeler, E. A., Baas, P., Gasson, P. E.** (1989). *IAWA list of microscopic features for hardwood identification* (Vol. 10). IAWA Leiden. <https://www.academia.edu/download/42652264/IAWA.Hardwood.List.pdf>
- Wheeler, E. A., Lehman, T. M., Gasson, P. E.** (1994). Javelinoxylon, an Upper Cretaceous Dicotyledonous Tree from Big Bend National Park, Texas, with Presumed Malvacean Affinities. *American Journal of Botany*, 81(6), 703-710. <https://doi.org/10.2307/2445648>
- Woodcock, D. W., Meyer, H. W., Prado, Y.** (2017). The Piedra Chamana fossil woods (Eocene, Peru). *IAWA Journal*, 38, 313-365. <https://doi.org/10.1163/22941932-20170175>