

Supporting Online Material for

Effects of Rapid Global Warming at the Paleocene-Eocene Boundary on Neotropical Vegetation

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This PDF file includes:

Materials and Methods
Figs. S1 to S14
Tables S1 and S5 to S20
References

Other Supporting Online Material for this manuscript includes the following: available at www.sciencemag.org/cgi/content/full/330/6006/957/DC1

Tables S1 to S4 as a zipped file

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Supplementary Online Material

Includes methods & expanded results, 14 supplementary figures, and 10 supplementary tables.

1. STATISTICS	2
2. STUDY SITES AND SAMPLING SUMMARY.....	2
3. GEOLOGY	2
3.1 REGIONAL GEOLOGY	2
3.2. LITHOLOGICAL DESCRIPTION OF THE SECTIONS	3
3.2.1 Mar 2x	3
3.2.2 Riecioto Mache	4
3.2.3 Gonzales	4
4. IDENTIFYING THE PETM INTERVAL	5
4.1. RADIOMETRIC DATING (U-PB ZIRCON GEOCHRONOLOGY)	6
4.1.1 U-Pb LA-ICPMS.....	7
4.1.2 U-Pb CA-TIMS	9
4.2. BIOSTRATIGRAPHIC DATUMS	10
4.3 STABLE CARBON ISOTOPES	12
4.3.1 Bulk Carbon Isotopic Analysis	12
4.3.2. Plant Biomarkers.....	13
5. DIVERSITY ANALYSES	14
5.1 STANDING DIVERSITY	15
5.2 PIECEWISE ANALYSIS.....	15
5.3 DCA ANALYSIS	16
5.4 CLUSTER ANALYSIS	16
5.5 WITHIN-SAMPLE DIVERSITY (RAREFACTION).....	17
5.6 SIMPSON INDEX	17
5.7 ANGIOSPERMS AND FERNS	17
5.8 FLORISTIC AFFINITIES	18
5.9 ARIDITY VS RAINFALL ANALYSIS	21
5.10 RATES OF ORIGINATION AND EXTINCTION	21
6. TEMPERATURE ANALYSIS.....	22
6.1 ESTIMATION OF TEMPERATURE.....	22
6.2 TEX ₈₆ ANALYSIS	23
6.3 CORE P2, NE COLOMBIA.....	24
6.3.1 Description Core P2, NE Colombia.....	24
6.3.2 Biostratigraphic Dating Core P2, NE Colombia	24
SUPPLEMENTARY REFERENCES	27

SUPPLEMENTARY FIGURES	30
SUPPLEMENTARY TABLES.....	32

1. Statistics

All the analyses, unless noted, were performed using *R for Statistical Computing* (S1), and the packages *Stratigraph* (S2), *Vegan* (S3) and *Cluster* (S4). All comparisons are the result of two-sided t-tests to evaluate the equality of means in two unpaired samples. *p* is reported for each test at the appropriate point in the text along with degrees of freedom (*df*) calculated using the Welch modification to account for different variances in the groups being compared.

2. Study Sites and sampling summary

Three sites were studied: Mar 2X, a 450-m-thick sediment core drilled in western Venezuela (11.0601° N, 72.1725° W, paleolatitude = 9.1° N); Riecito Mache, a 600-m-thick outcrop section in western Venezuela (10.7407° N, 72.3664° W, paleolatitude = 8.9° N), and Gonzales, a 170-m-thick outcrop section combined with a 600 m well in northeastern Colombia (8.28° N, 72.57° W, paleolatitude = 6.3° N). The study encompassed 489 carbon isotope samples (Table S1), and 357 palynological samples, recording 37,952 individual occurrences and 1104 morphospecies (Tables S1, S2 and S3). The Mar 2X Core comprised 165 palynological samples, 818 morphospecies and 25,281 occurrences (Table S2); the Riecito Mache section comprised 51 palynological samples, 262 species and 6,069 occurrences (Table S3); and the Gonzales section comprised 141 palynological samples, 244 species and 6,602 occurrences (Table S4). The Mar 2X Core has the highest palynological sample resolution, whereas the other two sites were used to confirm the patterns observed in Mar 2X. The Gonzales outcrop section is from a basin 320 km south of Mar 2x and the Riecito Mache outcrop section is a nearby site, 42 km southwest of Mar 2x.

3. Geology

3.1 Regional Geology

Basin geometries during the Paleogene in Colombia and Venezuela are very complex and do not seem to fit a single model. Studied sections come from two basins:

Catatumbo (Gonzales) and western Maracaibo (Rieci Mache and Mar 2x). Both basins have complicated subsidence histories due to eustatic and tectonic regimes that changed through time.

Catatumbo basin (Santander and craton-derived sediments). The basin sediments were derived from the Santander Massif, Merida Arch and the South American craton; the depositional systems prograde toward the Maracaibo Gulf (S5,S6). The regional progradation during the late Maastrichtian to late Paleocene is associated with the uplift of western massifs. A drop of base level due to a decrease in tectonic subsidence produced a sequence boundary during the late early Eocene, followed by coarse-grained fluvial deposition during the middle Eocene. Then, uplift of nearby western blocks led to the filling of a foredeep basin during the late Eocene to early Oligocene. Paleogene sediments are fully fluvial and there is no evidence of major time hiatuses (S5,S7,S8,S9).

Westernmost Maracaibo basin. The Paleocene-Lower Eocene succession in this area changes upsection, from carbonates with terrigenous material of the Guasare Formation to coal-bearing mudstone and sandstones of the Marcelina Formation. Overlying it are sandstones, mudstones, and conglomerate beds of the Misoa and Mostrencos Formations. In some areas toward the east of Maracaibo, the Misoa rests unconformably on top of the Marcelina, whereas there is no hiatus to the west (S5,S10,S11). This succession records syn-orogenic filling of a shallow marine basin that has progradational clastic wedge with source areas located to the west. The thin succession reported in this paper is at the lower segment of the Mostrencos Formation (Rieci Mache) and the middle segment of the Marcelina Formation (Mar 2x).

3.2. Lithological Description of the Sections

The PETM was identified at the middle Marcelina Formation in Mar 2x, at the lower Mostrencos Formation in Rieci Mache and at the upper Cuervos Formation in Gonzales. The Marcelina and lower Mostrencos Formations were accumulated in coastal plain deposits, whereas the Cuervos Formation was accumulated in fluvial deposits.

3.2.1 Mar 2x

The studied interval of the Mar 2x core (Fig. S1) consists of the middle part of the Marcelina Formation that contains the PETM. The sequence is characterized by cross-

bedded sandstone with shale and coal interbeddings, forming fining-upward intervals, up to 20 m-thick. This unit is interpreted as coastal and delta plain channels and associated overbank environments. Towards the top of the core (upper Marcelina) intercalated shale and ripple cross-laminated and bioturbated sandstone are more common, recording coastal and interdistributary bay and delta front environments. Lithological description was complemented by information provided by Petroleos de Venezuela-PDVSA (S12).

3.2.2 Riecito Mache

The sequence measured in the Riecito Mache section (Figs. S2-3) corresponds to the Paso Diablo and Mostrencos Formations. The Paso Diablo Formation consists of shale and coal with lenticular and tabular sandstone interbeddings, and cross-bedded sandstone with an erosive base forming fining-upward intervals up to 4 m thick. This unit was deposited in delta plain environments. The Mostrencos Formation contains cross-bedded sandstone with conglomeratic lenses, forming fining-upward intervals up to 15-m-thick, intercalated with shale and coal, and locally with tuffaceous sandstone. This unit is interpreted to represent low-sinuosity coastal plain environments. The PETM lies within the lower Mostrencos Formation. The section described herein was combined with a description and sampling of the same section conducted in the 1950's by H.W. Steckhoven (Royal Shell). Samples from the Steckhoven column, which were stored along with the stratigraphic column at the Core Library of PDVSA in Maracaibo, were analyzed by Rull (S11). His work showed a gradual increase in diversity from the Paleocene into the early Eocene. However, his study was based only on key biostratigraphic taxa, thus excluding a large number of taxa. Even with this biased data collection, Rull's analysis reached the same conclusion as ours, i.e. there is a gradual transition in the vegetation and increasing diversity in the Early Eocene.

3.2.3 Gonzales

An outcrop section and a well drilled at the outcrop section were used in this study (Figs. S4 and S5). The correlation of the outcrop section and the well is straightforward (see correlation points in Fig. S5). All samples for the analysis were expressed in units of the Gonzales well, because it had a more complete section. The upper segment of the Cuervos Formation, the segment containing the PETM, is

characterized by fine-grained successions interbedded with meter-scale channel-fill sandstones that record fluvial deposition in a basin with high subsidence.

4. Identifying the PETM interval

We have three independent lines of evidences indicating the PETM interval in the different sections.

1) Radiometric dating. Within the upper part of the PETM interval in Rieci Mache, a radiometric age of 56.09 ± 0.03 Ma was determined using a high-resolution technique (Chemical Abrasion– Thermal Ionization Mass Spectrometry). This age is within one of the possible ages estimated for the PETM (S13); given a duration of ~ 200 K for the PETM, its age would span from 56.33 to 56.13.

2) Biostratigraphic dating. Palynostratigraphy has been used in the region over the last 50 years as the main biostratigraphic tool by numerous oil companies and researchers. There are several taxa indicators for both the onset of the Eocene and the end of the Paleocene. Within the PETM interval, there are 10 key Eocene taxa originating and 3 key Paleocene taxa becoming extinct. This co-occurrence of events happens in all three sections (Mar 2x, Rieci Mache, and Gonzales). In summary, there is a strong biostratigraphic support for the PETM position.

3) The Carbon Isotope record, both in Bulk and compound specific carbon isotope records. Bulk isotope records of all cores indicate that PETM interval samples have a statistically lower bulk $\delta^{13}\text{C}$ values than late Paleocene samples (1.7 to $1.9\text{\textperthousand}$). The bulk carbon isotope records from our sections are noisier as compared to carbon isotope record obtained from carbonates of deep-water sediments, likely because several factors, other than the atmospheric values of $\delta^{13}\text{C}$, influence the $\delta^{13}\text{C}$ signal in a fluvial setting (S14). However, in spite of all possible biases, PETM interval has a statistically significant lower bulk $\delta^{13}\text{C}$ values than late Paleocene samples ($1.7\text{\textperthousand}$). This difference is the same value ($1.7\text{\textperthousand}$) of the difference between the mean value of PETM samples versus late Paleocene samples that is found in Wyoming (S15), the best studied PETM terrestrial site in the world. When the most negative sample found within the PETM is compared with the Late Paleocene (e.g., the peak of the carbon excursion), the PETM excursion in our sites (2.9 to $3.2\text{\textperthousand}$) is lower than in Wyoming ($3.8\text{\textperthousand}$) (S15), that could indicate that we

are missing the peak of the excursion. We measured $\delta^{13}\text{C}$ on *n*-alkanes in the Mar2x core, which targets vascular plants material only. The results show that the *n*-alkanes confirm the bulk isotope records and show a negative carbon isotope excursion in all odd carbon numbered *n*-alkanes measured and ranges between 2-3‰.

In conclusion, these three independent lines of evidence indicate all sections contain the PETM interval. In the following headings (4.1, 4.2 and 4.3), we expand on the explanation of each line of evidence.

4.1. Radiometric Dating (U-Pb Zircon Geochronology)

A first order dating of the PETM worldwide has remained elusive due to the absence of associated volcanic material. The PETM was dated by establishing its position relative to dated ash -17 that lies within magnetochron C24r (S16).

In Rieci Mache an intercalated felsic pyroclastic tuff was dated using TIMS (Chemical Abrasion– Thermal Ionization Mass Spectrometry) to 56.09 ± 0.03 Ma. This volcanism is linked to the northern Andes and Circum-Caribbean Paleogene magmatic province (S17,S18). The volcanic layer was found within the PETM, at meter 1290, which is 11 m below the top the PETM (at 1301 m), and 67 meters above the bottom of the PETM (at 1223 m). The age of the onset of the PETM can be estimated with our new date if sedimentation is assumed constant during this interval and the duration is between 170 kyr (S19) and 220 kyr (S20). The PETM onset is estimated at 56.24 Ma (assuming 170 kyr) to 56.28 Ma (assuming 220 kyr). This range is closest to the orbital cyclostratigraphy option 3 (56.33 Ma) for the age of the PETM onset (S16).

Due to the common existence of older zircon crystals within explosive volcanics, either as reworked grains from previous volcanic phases or as inherited zircons associated with melting of older crust (S21,S22), two U-Pb geochronological methods were used to constrain the age of the youngest zircon in the Rieci Mache tuff, which is interpreted as having formed during eruption. Zircons were mounted in epoxy, polished, and examined by cathodoluminescence imaging to reveal the internal structure. Only grains with a simple growth history that appeared to indicate a single-stage crystallization were analyzed. Forty-eight grains in the epoxy mount were analyzed using LA-ICP-MS (laser ablation inductively coupled plasma mass spectrometry) at Washington State University.

Thirty-two grains were identified as being part of the youngest population, ten of which were removed from the epoxy mount and analyzed by the more precise CA-TIMS (chemical abrasion thermal ionization mass spectrometry) method at Boise State University.

Due to the greater precision, we use the CA-TIMS results for our age interpretations. The $^{206}\text{Pb}/^{238}\text{U}$ dates from the six youngest grains are equivalent with a weighted mean of 56.09 ± 0.03 Ma (MSWD = 1.5, Fig. S6, Table S5, calculated with Isoplot 3.0 (S23)), which is interpreted as the eruption age of the Rieci Mache tuff. The error is the internal error based on analytical uncertainties only, including counting statistics, instrumental fractionation, subtraction of tracer solution, and blank and initial common Pb subtraction. It is the 2σ error expanded by the square root of the MSWD and the Student's T multiplier of n-1 degrees of freedom. This error should be considered when comparing our date with $^{206}\text{Pb}/^{238}\text{U}$ dates obtained by other laboratories with tracer solutions that were similarly calibrated using EARTHTIME gravimetric standards. When comparing our date with those derived from other decay schemes (e.g., $^{40}\text{Ar}/^{39}\text{Ar}$, $^{187}\text{Re}-^{187}\text{Os}$), systematic uncertainties in the tracer calibration and ^{238}U decay constant (S24) should be added to the internal error in quadrature. This error is ± 0.13 Ma. Errors on the individual analyses are 2σ internal errors based on analytical uncertainties only.

Three other grains yielded slightly older CA-TIMS $^{206}\text{Pb}/^{238}\text{U}$ dates of 56.39 ± 0.04 to 56.17 ± 0.04 Ma, and one other grain yielded a significantly older date of 67.5 ± 0.1 Ma. The older analyses are interpreted as being from grains that had prolonged magma residence or contained inherited cores

4.1.1 U-Pb LA-ICPMS

Heavy mineral concentrates of the <350 μm fraction were separated magnetically. Inclusion-free zircons from the non-magnetic fraction were then handpicked under a binocular microscope. Fifty zircons were mounted in epoxy and polished to half thickness for laser ablation analyses, using laser ablation–inductively coupled plasma–mass spectrometry. All LA-ICP-MS U-Pb analyses were conducted at Washington State University using a New Wave Nd:YAG UV 213-nm laser coupled to a ThermoFinnigan Element 2 single collector, double-focusing, magnetic sector ICP-MS. Operating

procedures and parameters are discussed in greater depth by Chang et al. (S25) and are only briefly outlined here. Laser spot size and repetition rate were 30 nm and 10 Hz, respectively. He and Ar carrier gases delivered the sample aerosol to the plasma. Each analysis consisted of a short blank analysis followed by 300 sweeps through masses 204, 206, 207, 208, 232, 235, and 238, taking approximately 35 seconds.

LA-ICP-MS isotopic analyses are affected by two forms of inter-element fractionation that must be corrected (S26). Time-dependent fractionation results from the more efficient volatilization of Pb over U as the laser excavates successively deeper levels in the ablation pit during an analysis, which in turn leads to an increase in $^{206}\text{Pb}/^{238}\text{U}$ and $^{207}\text{Pb}/^{235}\text{U}$ ratios with time (S27). It has been demonstrated that U/Pb fractionation is approximately linear over the short time interval of the analysis (S26). By definition, time-dependent fractionation is zero at the beginning of the analysis. Regression of time series data to the intercept at $t = 0$, therefore, yields the point at which time-dependent fractionation equals zero.

Time-independent (or static) fractionation is the largest source of uncertainty in LA-ICP-MS U-Pb geochronology and results from mass and elemental static fractionation in the plasma and also poorly understood laser-matrix effects (S26). The time-independent fractionation corrected by normalizing U/Pb and Pb/Pb ratios of the unknowns to the zircon standards (S25). For this study we used two zircon standards: Peixe, with an age of 564 Ma (S28), and FC-1, with an age of 1099 Ma (S29). Peixe was used to correct the $^{238}\text{U}/^{206}\text{Pb}$ and $^{235}\text{U}/^{207}\text{Pb}$ ratios and FC-1 was used to correct the $^{207}\text{Pb}/^{206}\text{Pb}$ ratios.

Common Pb can represent a proportionally large contribution to the total Pb in Mesozoic and younger U-poor zircons. However, common Pb is typically not significant in LA-ICP-MS analyses, most likely because it is concentrated in cracks and inclusions, which can be avoided. When this is not possible, the influence of common Pb is easy to recognize on Tera-Wasserburg diagrams because analyses tend to line up on a steep linear trajectory that can be anchored at a reasonable $^{207}\text{Pb}/^{206}\text{Pb}$ common lead composition (y-intercept) (S30). Common Pb corrections were made on these analyses using the ^{207}Pb method (S31). The U-Pb age was calculated using Isoplot (S23). The final crystallization ages that we report are using the algorithm of TUFFZIRC. Analyses

that are statistically excluded from the main cluster are shown in gray on the figure. The ages that we report in Table S6 are at the one sigma level and report only analytical error. The final age error (Table S6) was calculated using two uncertainties: the first is derived from the uncertainty of the TUFFZIRC age calculation alone, and the second represents the systematic uncertainty during that session (~1.1 %). The age uncertainty is determined as the quadratic sum of the TUFFZIRC error plus the total systematic error for the set of analyses (~1.2%). The obtained TUFFZIRC age of 55.8 ± 0.7 overlaps within error with the CA-TIMS age (Figure S7).

4.1.2 U-Pb CA-TIMS

From the major youngest population of zircon that were identified during LA-ICPMS analysis, which we interpreted as having formed during eruption of the tuff, ten single zircon analyses were selected for chemical abrasion–thermal ionization mass spectrometry (CA-TIMS) at Boise State University.

Zircons were subjected to a modified version of the chemical abrasion method of Mattinson (S32), reflecting analysis of single grains. Grains were removed from epoxy mounts after LA-ICPMS dating and cathodoluminescence imaging, and then placed in a muffle furnace at 900°C for 60 hours in quartz beakers. Single grains were then transferred to 3 ml Teflon PFA beakers and loaded into 300 μl Teflon PFA microcapsules. Fifteen microcapsules were placed in a large-capacity Parr vessel, and the crystals were partially dissolved in 120 μl of 29 M HF for 12 hours at 180°C. The contents of each microcapsule were returned to 3 ml Teflon PFA beakers, the HF removed and the residual grains were immersed in 3.5 M HNO₃, ultrasonically cleaned for an hour, and fluxed on a hotplate at 80°C for an hour. The HNO₃ was removed and the grains were rinsed twice in ultrapure H₂O before being reloaded into the same 300 μl Teflon PFA microcapsules (rinsed and fluxed in 6 M HCl during sonication and washing of the grains) and spiked with the Boise State University mixed ²³³U-²³⁵U-²⁰⁵Pb tracer solution. These chemically abraded grains were dissolved in Parr vessels in 120 μl of 29 M HF with a trace of 3.5 M HNO₃ at 220°C for 48 hours, dried to fluorides, and then re-dissolved in 6 M HCl at 180°C overnight. U and Pb were separated from the zircon

matrix using an HCl-based anion-exchange chromatographic procedure (S33), eluted together and dried with 2 μ l of 0.05 N H_3PO_4 .

Pb and U were loaded on a single outgassed Re filament in 5 μ l of a silica-gel/phosphoric acid mixture (S34), and U and Pb isotopic measurements made on a GV Isoprobe-T multicollector thermal ionization mass spectrometer equipped with an ion-counting Daly detector. Pb isotopes were measured by peak-jumping all isotopes on the Daly detector for 100 to 160 cycles, and corrected for $0.15 \pm 0.06\%$ /a.m.u. (2σ) mass fractionation. Transitory isobaric interferences due to high-molecular weight organics, particularly on ^{204}Pb and ^{207}Pb , disappeared within approximately 30 cycles, ionization efficiency averaged 104 cps/pg of each Pb isotope. Linearity (to $\geq 1.4 \times 10^6$ cps) and the associated deadtime correction of the Daly detector were monitored by repeated analyses of NBS982, and have been constant since installation. Uranium was analyzed as UO^{2+} ions in static Faraday mode on 10^{11} ohm resistors for 200 to 250 cycles, and corrected for isobaric interference of $^{233}U^{18}O^{16}O$ on $^{235}U^{16}O^{16}O$ with an $^{18}O/^{16}O$ of 0.00206. Ionization efficiency averaged 20 mV/ng of each U isotope. U mass fractionation was corrected using the known $^{233}U/^{235}U$ ratio of the tracer solution.

U-Pb dates and uncertainties were calculated using the algorithms of Schmitz and Schoene (S35), $^{235}U/^{205}Pb$ of 77.93 and $^{233}U/^{235}U$ of 1.007066 for the Boise State University tracer solution, and U decay constants recommended by Jaffey et al. (S24). $^{206}Pb/^{238}U$ ratios and dates were corrected for initial ^{230}Th disequilibrium using a Th/U[magma] = 3 using the algorithms of Crowley et al. (S36), resulting in an increase in the $^{206}Pb/^{238}U$ dates of ~0.09 Ma. All common Pb in analyses was attributed to laboratory blank and subtracted based on the measured laboratory Pb isotopic composition and associated uncertainty. U blanks are difficult to precisely measure, but are estimated at 0.07 pg.

4.2. Biostratigraphic Datums

Palynology has been used extensively as a biostratigraphic tool in Colombia and Venezuela over the past 50 years, mainly by the oil industry, with excellent results (S37,S38,S39,S40,S41,S42,S43). In Mar 2x, within the PETM, there is the last appearance datum (LAD) of three key Paleocene markers (S38,S39,S42,S44),

Foveotricolpites perforatus (2310.6 m or 7580.6 ft), *Bombacacidites annae* (2287.6 m or 7505.3 ft), and *Retidiporites magdalenensis* (2310.6 m or 7580.6 ft), as well as the first appearance datum (FAD) of several Eocene markers (S38,S39,S42,S44) including *Cyclusphaera scabrata* (2314.7 m or 7594.4 ft), *Rhoipites hispidus* (2308.9 m or 7575.2 ft), *Retitrescolpites? irregularis* (2316.8 m or 7601.3 ft), *Tetracolporopollenites transversalis* (2315.5 m or 7596.9 ft), *Margocolporites vanwijhei* (2287.6 m or 7505.3 ft), *Tetracolporopollenites maculosus* (2294.2 m or 7527 ft), *Retibrevitricolpites triangulatus* (2316.8 m or 7601.3 ft), *Racemonocolpites facilis* (2311.1 m or 7582.5 ft), *Corsinipollenites undulatus* (2294.2 m or 7527 ft), and *Psilastephanocolporites fissilis* (2311.1 m or 7582.5 ft). Depths are given in both meters and feet, as the Mar2x core was taken using the English system, commonly used in the oil industry.

Biostratigraphy also confirms the position of the PETM in Rieci Mache (the LAD of *Foveotricolpites perforatus* 1261 m, *Bombacacidites annae* 1261 m, *Retidiporites magdalenensis* 1232 m, and the FAD of *Cyclusphaera scabrata* 1229.75 m, *Retitrescolpites? irregularis* 1229.75 m, *Tetracolporopollenites transversalis* 1229.75 m, *Margocolporites vanwijhei* 1295.45 m, *Retibrevitricolpites triangulatus* 1229.75 m, *Striatopolis catatumbus* 1229.78 m, *Ranunculacidites operculatus* 1295.45 m, *Rhoipites guianensis* 1281.8 m). Biostratigraphy also confirms the PETM in Gonzales (LAD of *Bombacacidites annae* (246.8 m or 810ft) and the FAD of *Cyclusphaera scabrata* (227.8 m or 747.4 ft), *Rhoipites hispidus* (226.3 m or 742.5 ft), *Racemonocolpites facilis* (223.5m or 733.5 ft), and *Rhoipites guianensis* (227.35 m or 745.9 ft)).

The Paleocene/Eocene boundary is defined by the Carbon Isotope Excursion (CIE), which lies within the lower part of planktonic foraminifera zone P5 (*Morozovella velascoensis*) of Berggren et al. 2005 (S45,S46). Dating terrestrial strata in tropical South America traditionally has been done with palynology (S38,S40,S42,S47,S48,S49). The calibration of these palynological zones to the geological time scale has been developed over many years and it is not an easy task. Researchers, mostly from the oil industry, have been able to analyze marine cores that have terrestrial palynomorphs, in order to calibrate the palynological zones (S38,S42). Additionally, the long-term record of $\delta^{13}\text{C}$ has been used as a calibration tool (S50,S51). The upper part of palynological zone *Retidiporites magdalenensis* of Germenaad et al. (S38), which contains the taxa used here

to indicate the late Paleocene (as described above) has been calibrated to planktonic foraminifera zone P4 of Berggren et al. (S45) by the co-occurrence of *Globorotalia pseudomenardii* and *Morozovella velascoensis* (S38). Zone P4 is dated as late Paleocene (S46). Palynological zone *Retibrevitricolporites triangulatus* of Germenaad et al. (S38), which contains the taxa used here to indicate the early Eocene (as described above), has been calibrated to planktonic foraminifera zone P5 of Berggren (S45) by the occurrence of *Morozovella velascoensis* and the absence of *Globorotalia pseudomenardii* (S38). Zone P5 as been dated as early Eocene to latest Paleocene (S46). The comparison of the long-term global $\delta^{13}\text{C}$ record of Zachos et al. (S52) to the Maastrichtian to middle Eocene record of the $\delta^{13}\text{C}$ in strata from Colombia has also been used to calibrate the palynological zones of the Paleogene of Colombia (S50,S51). The exact relationship of the palynological zones with the Paleocene/Eocene boundary (the PETM) has never been studied in detail. In this study we found that the transition from the *Retidiporites magdalenensis* to the *Retibrevitricolporites triangulatus* zone happens within the PETM interval, as confirmed by the radiometric dating.

4.3 Stable carbon isotopes

4.3.1 Bulk Carbon Isotopic Analysis

Stable carbon-isotope values of bulk organic matter ($\delta^{13}\text{CTOM}$) were measured via flash-pyrolysis at 1100 °C in a Costech elemental analyzer fitted to a Thermo Finnigan Delta plusXP isotope ratio mass spectrometer (Department of Geological Sciences at Indiana University-Bloomington). Carbonate present in the samples was removed by HCl digestion. Analytical precision and accuracy were determined on the basis of repeated analysis of two internal lab standards calibrated against the internationally accepted V-PDB standard. Overall uncertainty was better than 0.08‰. Organic carbon content (TOC) was determined on the basis of the liberated CO₂ in the elemental analyzer. $\delta^{13}\text{C}$ values are given in Table S1.

Bulk organic $\delta^{13}\text{C}$ values of Mar 2x were also complemented by a Middle to Late Paleocene record of $\delta^{13}\text{C}$ in a nearby core, 58 km to the west (S53). The PETM was identified in the 2287–2322 m interval. $\delta^{13}\text{C}$ values of the PETM are significantly more negative than that of the Middle-Late Paleocene (mean -27.3‰ vs. -25.6‰, $p < 0.0002$,

df: 104.556), and slightly more negative than the $\delta^{13}\text{C}$ of the earliest Eocene (-27.1), although the difference is not significant ($p < 0.193$, df: 110.82). At Rieci Mache, the PETM is contained within 1223-1301 m interval [PETM -27.4, Paleocene -25.7, $p < 0.001$, df: 7.799; Early Eocene -26.2, $p < 0.005$, df: 7.824]. At Gonzales, the PETM is contained within 222.5- 251.5 m interval [PETM -27.4‰, Paleocene -25.5‰, $p < 0.001$, df: 32.849; Early Eocene -24.9‰, $p < 0.001$, df: 25.473]. The most negative isotope value within the PETM compared to late Paleocene values (e.g., the peak of isotope excursion) is 2.92‰ at 2293.6 m in Mar 2x, 3.2‰ at 1231.5 m in Rieci Mache, and 3.04‰ at 227.3 m in Gonzales.

Because $\delta^{13}\text{C}$ values of bulk sediments can be affected by the total organic carbon of a sample (S15); we applied the Wing residuals method (S15) to the $\delta^{13}\text{C}$ record of the three sites. The resulting pattern does not differ significantly from the raw data (Figs. S8, S9, S10).

4.3.2. Plant Biomarkers

Sediments were extracted with 2:1 (v/v) dichloromethane: methanol using an accelerated solvent extractor (ASE 300; Dionex Corporation) at 120°C, 1,500 p.s.i., for 25 minutes. Lipid fractions were separated by silica gel column chromatography using an elution sequence of hexane, dichloromethane, and methanol. Cyclic and branched alkanes were separated from normal and isoalkanes by urea adduction. The hydrocarbon fraction (hexane fraction) was dried under a stream of N₂ and dissolved in a mixture of methanol-saturated urea, pentane and acetone (200ul each). The resulting urea crystals were extracted with hexane yielding cyclic/branched alkanes. Remaining crystals are dissolved in 1:1 (v/v) H₂O: methanol, then extracted with hexane to yield the *n*-alkane fraction. We also performed silver nitrate impregnated silicagel chromatography as an additional clean-up step before compound specific isotopic analysis. For *n*-alkane analysis, low values of Carbon Preference Index (CPI; $\sum(\text{C}_{25}+\text{C}_{27}+\text{C}_{29}+\text{C}_{31})/\sum(\text{C}_{24}+\text{C}_{26}+\text{C}_{28}+\text{C}_{30})$) were used to eliminate samples that could be biased by contributions from other marine sources such as aquatic plants. All samples with a CPI value less than 1.38 were not used for isotopic analysis.

The *n*-alkane fraction was analyzed for stable carbon and hydrogen isotopic

compositions on a Thermo Finnigan MAT 253 mass spectrometer interfaced with a Thermo Finnigan Trace GC Combustion III (for carbon) and High Temperature Conversion (for hydrogen) systems. A J&W Scientific DB-1 capillary column was used to separate individual n-alkanes. Temperature was programmed from 60°C (held for 1 min) at 6°C min⁻¹ to 320°C and held for 25 min isothermally. A programmed temperature vaporizing injector was used with Helium as a carrier gas with a column flow rate of 2.0 ml min⁻¹. Carbon isotopic compositions are expressed relative to VPDB standard. The analytical accuracy and precision are determined based on repeated analysis of a standard n-alkane mixture (MixB2 containing C16-C30 n-alkanes; isotopic ratios measured offline by A. Schimmelmann, Biogeochemical Laboratories, Indiana University). The standard error of n-alkane δ¹³C (based on duplicates of n samples) was ±0.8‰ or better. H₂ was obtained from organic hydrogen by pyrolytic conversion at 1400°C. The H₃⁺ factor (proportionality constant between the concentration of H₂ and H₃⁺) was determined daily using reference H₂ gas. Analytical accuracy and precision of the system were determined using the mixture of n-alkane compounds. Standard error was generally less than ±5‰. Results are summarized in Table S7, Figure S11 (C n-alkane), and Figure S12 (D n-alkane).

5. Diversity Analyses

Pollen and spore morphotypes reflect mostly generic diversity (S38) with some few types reflecting diversity at the species and family levels, although the natural affinity of many of the morphotypes in our database is still unknown. However, there is no reason to expect a systematic bias in the taxonomic representation of the pollen or spores that would affect any segment of the diversity pattern. Palynological data have been previously used to study plant diversity over different time scales (S54,S55,S56,S57,S58,S59,S60,S61).

Species morphologies can be seen in our online morphological database at <http://biogeodb.stri.si.edu/jaramillo/palynomorph/>. An additional 390 Mb Filemaker file with photos and descriptions can be downloaded from the Smithsonian Tropical Research Institute Data Repository at <http://biogeodb.stri.si.edu/jaramillo/paper/paperData.html>. The raw abundance counts for each of the sites are given in Tables S2, S3, and S4.

5.1 Standing Diversity

We used the range-through method to calculate standing diversity, because it decreases the bias produced by changes in facies and depositional environments (Table S8). All species with single occurrences were eliminated from the standing diversity analysis. Standing diversity was not calculated for Gonzales because samples of the PETM interval had higher counts than both Eocene and Paleocene samples (counts of 111.1 vs. 259.6, $p < 0.001$), and higher sample counts artificially increase standing diversity.

5.2 Piecewise Analysis

The edge effect (S62) in the standing diversity curve was estimated using a piecewise regression. This regression assumes that there are two different regression functions to the same data (S63) and attempts a two-segment fit of the data. The breakpoint is the intersection of the two fitted regression lines. The regression iteratively tries all possible positions of the breakpoint and chooses the one that produces the lowest residual sum of squares (S64). The model to fit follows the algorithm described by Duggleby and Ward (S65) for a two-segment linear regression: $y = y_T + [(m_L + m_R)(x - x_T) - (m_L - m_R) |x - x_T|] / 2$ $y = \text{FAD}$ or LAD , $x = \text{species}$, $x_T = \text{breakpoint species}$, $y_T = \text{breakpoint FAD or LAD}$, $m_L = \text{slope left of breakpoint}$, $m_R = \text{slope right of breakpoint}$. Following is the code used in R and the package *Stratigraph* to perform the analysis.

```
##FAD EDGE
mar2x.edge=strat.column(counts=t(mar2x.countsNoSingles),depths=mar2x.depthFeet,sample.labels=mar2x.Species)#Singles out
mar2x.FAD=fads(mar2x.edge)#FAD, singles excluded
mar2x.LAD=lads(mar2x.edge)#LAD, singles excluded

fad.edge<-mar2x.FAD[which(mar2x.FAD>7400)]##time restricted to edge effect for FAD, up to 64My
fad.edge<-sort(fad.edge,decreasing = TRUE)##fad arranged from oldest to youngest
span=length(fad.edge)#length of the analysis, to be included in the next line
cumuedge.fad<-c(1:span)
step1<-numeric(span)##first segment of piecewise
for (i in (1:span)){
```

```

step1[i]<-sum(resid(lm(fad.edge[1:i]~cumuedge.fad[1:i]))^2)
}
step2<-numeric(span)##last segment of piecewise
for (i in (1:span)){
  step2[i]<-sum(resid(lm(fad.edge[i:span]~cumuedge.fad[i:span]))^2)
}
piecewise<-step1+step2## sum of first and second segment
breakpoint=which(piecewise==min(piecewise))## the position of the minimum value,
data 148
fad.edge[breakpoint]##7698.1 Ma

##LAD EDGE
mar2x.edge=strat.column(counts=t(mar2x.countsNoSingles),depths=mar2x.depthFeet,sample.labels=mar2x.Species)
mar2x.FAD=fads(mar2x.edge)#FAD, singles excluded
mar2x.LAD=lads(mar2x.edge)#LAD, singles excluded

lad.edge<-mar2x.LAD[which(mar2x.LAD<7400)]##time restricted to edge effect for FAD, up to 64My
lad.edge<-sort(lad.edge)##lad arranged from youngest to oldest
span=length(lad.edge)#length of the analysis, to be included in the next line
cumuedge.lad<-c(1:span)
step1<-numeric(span)##first segment of piecewise
for (i in (1:span)){
  step1[i]<-sum(resid(lm(lad.edge[1:i]~cumuedge.lad[1:i]))^2)
}
step2<-numeric(span)##last segment of piecewise
for (i in (1:span)){
  step2[i]<-sum(resid(lm(lad.edge[i:span]~cumuedge.lad[i:span]))^2)
}
piecewise<-step1+step2## sum of first and second segment
breakpoint=which(piecewise==min(piecewise))## the position of the minimum value,
data 148
lad.edge[breakpoint]##6986.1

```

5.3 DCA Analysis

A Detrended Correspondence Analysis (DCA) (S66) was performed, using the function *decorana* from the package VEGAN. The DCA was performed on the composite section after the range-through assumption. Singletons were excluded.

5.4 Cluster Analysis

An agglomerative cluster analysis (S67), using Euclidean distance, was performed

using the function *agnes* from the package *Cluster*. The Cluster was performed on the composite section after the range-through assumption. Singletons were excluded. The agglomerative coefficient, which measures the amount of clustering structure found, was 0.8761647 for Mar 2x, 0.7975151 for Rieci Mache, and 0.7949479 for Gonzales (Fig. S13).

5.5 Within-Sample Diversity (Rarefaction)

The within-sample diversity was analyzed using rarefaction (S68). The number of morphotypes found at counts of 100, 120 and 150 grains was calculated for each sample, using the package VEGAN (Table S8). In Gonzales, at 100-counts, the Paleocene is lower than the PETM (12.5 vs. 18.1, $p < 0.06$, $df: 5.04$), and the PETM is lower than the Eocene (18.1 vs. 26.2, $p < 0.07$, $df: 7.123$). In Rieci Mache, rarefaction was not calculated because very few samples had counts greater than 100 grains.

5.6 Simpson Index

The Simpson Index (S69) was calculated using *Stratigraph*, and performed in samples with a count above 80 grains (Table S8). In Rieci Mache, the Paleocene (0.74) is slightly lower than the PETM (0.76), $p < 0.82$, $df: 7.9$. One PETM sample (depth 1223) has the lowest Simpson value (0.2), much lower than all other samples of either the Paleocene or the PETM, and it is almost fully dominated by the species *Proxapertites "minutihumbertoides"*. If this sample is removed from the comparison, the Paleocene (0.74) is lower than the PETM (0.81), although the difference is not significant ($p < 0.503$, $df: 5.9$). In Gonzales, the Paleocene (0.57) is lower than the PETM (0.79, $p < 0.001$, $df: 19.148$), and the PETM is lower than the early Eocene (0.88, $p < 0.016$, $df: 11.9$).

5.7 Angiosperms and Ferns

Only samples with counts greater than 80 grains are considered in this analysis. In Mar 2x, angiosperm relative abundance is lower in the Paleocene (76.8%) than in the PETM (84.6%; $p < 0.07$, $df: 37.7$), whereas it is similar in the PETM and Eocene (86.9%; $p < 0.28$, $df: 24.03$). Angiosperm standing diversity is lower in the Paleocene

(136.36) than in the PETM (165.1, $p < 0.001$, df: 28.34), which itself has lower diversity than the Eocene (190.4, $p < 0.001$, df: 25.06). Fern spore standing diversity, on the other hand, does not change from the Paleocene (37.2) to the PETM (37.5; $p < 0.74$, df: 35.5), but it does increase slightly in the Eocene (40.4) ($p < 0.0001$, df: 27.7). This pattern is also seen in Gonzales and Riecito Mache.

In Riecito Mache, relative abundance of spores does not significantly change across the PETM (16.7 in Paleocene, 8.4 in PETM, $p < 0.5317$, df: 4.794), nor does that of pollen (Paleocene = 83.2 vs. Eocene = 91.5, $p < 0.531$, df: 4.7). Spore standing diversity is lower in the Paleocene (19.5) than in the Eocene (22.3), $p < 0.015$, df: 19.7, as well as pollen standing diversity [Eocene (72.8). Paleocene (56.8), $p < 0.0001$, df: 14.7].

In Gonzales, spore relative abundance slightly decreases from the Paleocene (29.5) to the Eocene (5.8, $p < 0.0001$, df = 54.); angiosperm pollen also increases in abundance (71.8 vs. 94.9). Spore standing diversity (after range-through) does not change from the Paleocene to the Eocene (12.6 vs. 12.5, $p < 0.87$, df: 26.1). Angiosperm pollen became more diverse in the Eocene compared to the Paleocene (54.8 vs. 80.8, $p < 0.0001$, df: 23.4).

5.8 Floristic Affinities

Floristic affinities were determined from the following sources: (S9,S38,S40,S41,S47,S48,S49,S70,S71,S72,S73,S74,S75,S76,S77,S78). In the following list we specified the natural affinities of the taxa, grouped by Family, which we used in the analysis.

Arecaceae

Arecipites regio

Baculapollenites grimsdaloide

Bacumonocolpites sp.

Echimonocolpites sp.

Gemmamonocolpites (all species)

Gemmastephanocolpites (all species)

Mauritiidites (all species)

Monocolpopollenites (all species)

Monosulcites

Psilamonocolpites (all species)

Racemonocolpites (all species)

Spinizonocolpites (all species)
Trichotomosulcites (all species)

Olacaceae
Anacolosidites cf. *luteoides*

Bombacoideae
Bombacacidites (all species)
Retistephanocolporites minimus (Bombacaceae type)

Onagraceae
Corsinipollenites

Fabaceae
Crassiectoapertites
Margocolporites
Polyadopollenites
Striatopolis catatumbus

Euphorbiaceae
Croton type
Crototricolpites pachidermatus
Crototricolpites sp.1DO
Ranunculacidites operculatus
Retitrescolpites? *irregularis*

Ctenolophonaceae
Ctenolophonidites lisamae
Verrustephanocolpites rugulatus

Proteaceae
Echitriporites trianguliformis
Echitriporites trianguliformis orbicularis
Proteacidites
Retidiporites magdalenensis

Araceae
Ephedripites chomotrileticus
Ephedripites rizadus
Proxapertites extrañus
Proxapertites heterofoveolatus
Proxapertites cursus
Proxapertites minutus
Proxapertites operculatus
Proxapertites psilatus
Proxapertites sp.

Proxapertites verrucatus
Proxapertites imperialis
Proxapertites immensus
Proxapertites nixinatus
Spathiphyllum sp. 1_PETM
Spathiphyllum vanegensis

Annonaceae

Proxapertites aff. *tertiaria*
Proxapertites magnus
Proxapertites humbertoides
Proxapertites *tertiaria*
Proxapertites minutihumbertoides
Proxapertites heterofoveolatus
L. proxapertitoides proxapertitoides
L. proxapertitoides reticuloides
Longapertites (all species)

Ericaceae

Ericipites aff. *annulatus*

Moraceae

Momipites (all species)
Psiladiporites (all species)

Poaceae

Monoporopollenites (all species)

Polypodiaceae

Polypodiisporites

Convolvulaceae

Perfotricolpites (all species)

Rhizophoraceae

Paleosantalaceaepites (all species)
Zonocostites (all species)

Podocarpaceae

Podocarpidites

Sterculioideae

Rhoipites miniguianensis
Rhoipites guianensis

Sapotaceae

Tetracolporopollenites (all species)

Ulmaceae

Ulmoideipites krempii

Myrtaceae

Syncolporites poricostatus

Pelliceriaceae

Lanagiopollis (all species)

Passifloraceae

Spirosyncolpites spiralis

5.9 Aridity vs Rainfall Analysis

Plant families identified in this study were classified as wet or dry, according to the analysis of Punyasena (S79), that identified Family precipitation preferences based on the Gentry's 144-transect neotropical plant database (Table S9). Fabaceae are both abundant in high rainfall and dry habitats within the tropics (S80). However, to be more conservative, Fabaceae was considered to be an indicator of dryness. Then, the sum of abundances of the families corresponding to each category (wet versus dry) was calculated for each sample of the Mar2x core (only samples with sums larger than 80 grains were used). The results (Table S9) indicate that both Paleocene and Eocene samples are dominated by families indicating wet habitats and there is not a significant difference in their combined relative abundances across the Paleocene-Eocene (Paleocene= 64%, Eocene= 61%, t-test, $p < 0.49$, df: 32.5). Abundance of dry elements (e.g., grasses) represents <2% of the assemblage (Paleocene=0.7%, Eocene=2%).

5.10 Rates of Origination and Extinction

Per capita rates of origination and extinction were calculated following Foote (S62) using the package *Stratigraph*. The bin used was the stratigraphic thickness of the PETM in the Mar 2x site (which was assumed to have accumulated for 200,000 years, the upper estimate of the duration of the PETM (S81), in order to be able to compare the PETM interval with levels below and above. These rates were not calculated in either

Riecito Mache or Gonzales, because the analyzed stratigraphic interval was not long enough.

6. Temperature Analysis

6.1 Estimation of Temperature

We estimate the mean annual temperature (MAT) of the region during the Late Paleocene, using both published information and data measured in this study (TEX_{86} from P2 core, a nearby marine site, 300 km west of Mar2x, that is described in subsequent headings 6.1 and 6.2). Sea-surface temperature (SST) from Tanzania was $\sim 29^\circ\text{C}$ at the uppermost Paleocene (S82). Published temperatures for Cerrejon, a Late Paleocene terrestrial site that is 50 km west of Mar2x, ranges from 28 to 34°C (S15,S83,S84). Using a snake paleothermometer, Head et al (S79, S80) produced two estimates of MAT, first $30\text{-}34^\circ\text{C}$ (S83), and later a refined estimation of $28\text{-}31^\circ\text{C}$ (S84). Here, we use the cooler estimate ($28\text{-}31^\circ\text{C}$) that seems to be in agreement with other proxies.

Our own estimates for the Late Paleocene, derived from TEX_{86} of core P2, yielded TEX_{86} values that were relatively low in the Late Paleocene (0.65-0.73, Table S10). To estimate SST from TEX_{86} , two calibrations (S85,S86) were applied that yielded similar estimates with an average of $28.2 \pm 1.4^\circ\text{C}$ (S85) and $27.3 \pm 1.1^\circ\text{C}$ (S86) for this time period. In summary, considering all proxies together, the range of MAT for the late Paleocene in the region was likely $28\text{-}31^\circ\text{C}$.

We estimated the increase in MAT during the PETM, using both published data and also TEX_{86} data from P2 core. Zachos et al. (S87) found an increase of $3\text{-}5^\circ\text{C}$ during the PETM in tropical Pacific sea-surface temperature (SST), slightly lower than estimates from high and mid latitude sites (S88,S89). We do not have a direct estimation of the temperature of the PETM in any of our sites, as they did not contain suitable sediments. Instead, we used the SST of the Early Eocene that was determined in core P2, as a proxy for the MAT during the PETM, because the absolute temperature increase was similar in both the Early Eocene and the PETM. Early Eocene TEX_{86} values were higher than the Paleocene (0.75-0.87, Table S10). SST estimates are $31.5 \pm 2.3^\circ\text{C}$ (S85) and $29.7 \pm -1.6^\circ\text{C}$ (S86), or a $\sim 2\text{-}4^\circ\text{C}$ increase relative to late Paleocene values. This

increase in SST calculated by TEX_{86} is in agreement with the 3-5 °C calculated by Zachos et al. (S87). Therefore, a conservative estimate of the increase in MAT during the PETM could be in the order of 3°C assuming a similar increase in MAT compared to SST. If the MAT of the Late Paleocene was 28-31 °C, then the MAT during the PETM was likely ~31-34 °C.

6.2 TEX_{86} analysis

Samples from a sediment core from Northeastern Colombia (P2, 9.541679° N, 75.337792° W, paleolatitude= 7.06° N, 77.1° W, paleolatitude calculated using GPlates (S90)), were analyzed for TEX_{86} (Table S10). Organic compounds were extracted from powdered and freeze-dried sediments with dichloromethane (DCM)/ methanol (MeOH) (9:1, v/v) by using the accelerated solvent extraction technique (Dionex). Excess solvent was removed using a rotary evaporation with vacuum. The total extracts were separated in polar and apolar fractions over an activated Al_2O_3 column using hexane:DCM (1:1, v/v) and DCM:MeOH (1:1, v/v), respectively. The polar fraction was then dissolved in a 99:1 (v/v) hexane/isopropanol mixture, and sieved using a 0.45 μm , 4 mm diameter polytetrafluoroethylene (PTFE) filter, before being analyzed using a high-performance liquid chromatography/atmospheric pressure positive ion chemical ionization mass spectrometer (HPLC/ APCI-MS). HPLC/APCI-MS analyses were done according to Schouten et al. (S91) using an Agilent 1100 series LC/MSD SL and separation and a Prevail Cyano column (2.1 × 150 mm, 3 mm; Alltech), maintained at 30 °C. The GDGTs were eluted using a changing mixture of hexane and isopropanol as follows: 99% hexane to 1% propanol for 5 minutes, then a linear gradient to 1.8% isopropanol for 45 minutes. Flow rate was 0.2 ml per minute. Single ion monitoring was set to scan the 5 $[\text{M}+\text{H}]^+$ ions of the GDGTs with a dwell time of 237 ms for each ion. GDGT signals were generally low, especially at the lower levels where intense weathering was observed, suggesting that GDGTs were degraded in these sections. The values reported here were those where all GDGT isomers used in the TEX_{86} were above the limit of quantification (S92). This threshold resulted in the rejection of about two-thirds of the data of all sediments analyzed. We applied the core top calibration equations of Kim et al.(S85):

$$T = 38.6 + 68.4 \times \log(\text{TEX}_{86})$$

and that of Liu et al. (S86):

$$T = -16.33 \times 1/\text{TEX}_{86} + 50.475$$

In order to translate TEX_{86} values into an estimate of mean annual sea-surface temperature. The calibrations give roughly similar SST estimates for the late Paleocene but the Liu et al. (S86) calibration gives lower early Eocene estimates because it is non-linear toward higher temperatures. BIT values, a proxy for the input of soil organic matter, were below 0.2 in all cases, suggesting that the TEX_{86} values were not affected by a contribution of soil-derived tetraether lipids.

6.3 Core P2, NE Colombia

6.3.1 Description Core P2, NE Colombia

Core P2 (9.541679° N , $75.337792^\circ \text{ W}$) was drilled in the northeastern region of Colombia in the Sinu-San Jacinto. During the Paleocene–Eocene transition, the northeastern part of South America was under dextral transpression, with oblique subduction of the Caribbean Plate under continental crust of northern Colombia (S93). Modern paleoenvironmental models (S94) indicate the presence of a shallow platform and deltaic systems in the Colombian Caribbean region (Sinu-San Jacinto Fold Belt). Core P2 (Fig. S14) has massive dark-gray to black calcareous mudstones, rich in organic matter rich, occasionally with horizontal bioturbation or plane-parallel beds; this lithology has intercalation of fining-upward sequences of calcareous litarenites, generally massive, with horizontal bioturbation and wavy lamination, and flaser and hummocky structure. Sediments suggest deposition in a prodeltaic environment. There are no major lithological changes across the Paleocene–Eocene transition.

6.3.2 Biostratigraphic Dating Core P2, NE Colombia

The age of the core was determined by using the planktonic foraminiferal zonation of Berggren (Fig S14) (S45,S95). Pollen recovery in the core was scarce, but confirms the age given by foraminifera. The late Paleocene was recognized by planktonic foraminifera zones P3 and P4. Zone P3 (from the first appearance of *Morozovella angulata* and/or *Igorina pusilla* to the first appearance of *Globanomalina pseudomenardii*) is represented by an interval dominated by *Morozovella aequa*, *M. subbotinae* and *Subbotina variospira*. Nine samples from the 189.36-134.26 m interval

contain the P3 assemblage. However, this zone cannot be sharply defined. According to various sources (S45,S46,S95), Zones P3 and P4 are distinguished by the first appearance of *Morozovella angulata* and/or *Igorina pusilla* and the first appearance of *Globanomalina pseudomenardii*. In the core, *M. angulata* occurs only in two samples from the upper part of Zone P4; *I. pusilla* was not recognized. The first appearance of *Globanomalina pseudomenardii* is at 211.13m. Other species *Morozovella aequa* and *M. acuta* usually occurs slightly above the base of Zone P3. The benthic assemblage is dominated by *Spiroplectammina grzybowskii* and *Rzehakina epigona*. Zone P4 is defined as the total range of *Globanomalina pseudomenardii* (S45,S46,S95). *G. pseudomenardii* occurs in the interval from 211.13 to 309.62 m, whereas definite Zone P5 starts at 316.62 m, as discussed below. The absence of *G. pseudomenardii* in the 316.62–334m interval may be due either to poor preservation of planktonic foraminifers or to extinction of the taxon. The Zone P4 marker (*Globanomalina pseudomenardii*) is last observed at 309.62 m and the last appearance of *Morozovella acuta* is at 316.62 m. The last appearances of *Morozovella aequa* and *Globanomaina chapmani* are at 334 m. Four samples were studied between these last two levels. Among them, one (at 319.88m) has no planktonic foraminifers and calcareous nannofossils are also absent, and a stratigraphic designation is difficult. The sample at 316.62 m contains neither *G. pseudomenardii* nor *Morozovella subbotinae*. Because the P4/P5 zone boundary is primarily defined by either the last appearance of *G. pseudomenardii* or the first appearance of *Morozovella subbotinae*, the sample should belong to Zone P5. However, the biostratigraphic zonation of the 325.66–334 m interval is represented by a condensed interval treated as belonging to Zone P6, because it does not contain *M. acuta*. Because of poor preservation, the diversity of planktonic foraminiferal assemblages in Zone P5 is lower than in Zones P4 and P6. The early Eocene was recognized by zone P6. Species occurring in Zone P5 are common open-marine taxa. The original diversity may have been higher, because dissolution may have removed some fragile shells and recrystallization also makes recognition of some zones difficult. As stated above, Zone P6 cannot be unequivocally separated from Zone P5 on the basis of planktonic foraminiferal and nannofossils biostratigraphy alone because of the rare occurrence of micro and nannofossils between 316.88 and 334 m. As a result of rare occurrence secondary datum events to mark the definite top, Zone P5

includes the last appearance of *Morozovella acuta* and Zone P6 includes the last appearance of *Morozovella aequa* and *Globanomalina chapmani*. In addition, the calcareous nannofossils *Chiasmolithus bidens*, *Fasciculithus tympaniformis* and *Neochiastozygus saepes* occur in the base of nannofossil Zone NP10, at 325.66 m. Planktonic foraminiferal biostratigraphy places the base of Zone P6 and the base of the Eocene above the last appearance of *Morozovella acuta* at 316.62 m, as discussed above. Nannofossil biostratigraphy indicates an early Eocene age at 325.66 m.

Supplementary References

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Supplementary Figures

Fig. S1. Lithological description of the Mar 2x Core, Maracaibo, Venezuela.

Fig. S2. Polygonal of the Riecito Mache section, Maracaibo, Venezuela.

Fig. S3. Lithological description of the Riecito Mache, Maracaibo, Venezuela.

Fig. S4. Polygonal of the Gonzales section, Catatumbo, Colombia.

Fig. S5. Lithological description of the Gonzales outcrop and Gonzales well, Catatumbo, Colombia.

Fig S6. Plot of the ^{206}Pb - ^{238}U dates from the youngest zircon from the Riecito Mache tuff obtained by the U-Pb CA-TIMS method. Plotted with Isoplot 3.0 (S23). Error bars are 2σ internal errors. Weighted mean date is represented by the grey box behind the error bars. Error bars in white were not used in the weighted mean calculation.

Fig S7. U-Pb LA-ICP-MS TUFFZIRC age from the Riecito Mache tuff. Plotted with Isoplot 3.0(S23).

Fig. S8. Carbon isotope data ($\delta^{13}\text{C}$) vs. Stratigraphic position, Mar 2x section following the method by (S15). A. Total organic carbon (TOC) vs. carbon isotope data ($\delta^{13}\text{C}$). B. Residual of correlation TOC - $\delta^{13}\text{C}$ vs. the stratigraphic position. Pattern is very similar to raw $\delta^{13}\text{C}$ data.

Fig. S9. Carbon isotope data ($\delta^{13}\text{C}$) vs. stratigraphic position, Riecito Mache section following the method by (S15). A. Total organic carbon (TOC) vs. carbon isotope data ($\delta^{13}\text{C}$). B. Residual of correlation TOC - $\delta^{13}\text{C}$ vs. the stratigraphic position. Pattern is very similar to raw $\delta^{13}\text{C}$ data.

Fig. S10. Carbon isotope data ($\delta^{13}\text{C}$) vs. stratigraphic position, Gonzales section following the method by (S15). A. Total organic carbon (TOC) vs. carbon isotope data ($\delta^{13}\text{C}$). B. Residual of correlation TOC - $\delta^{13}\text{C}$ vs. the stratigraphic position. The pattern observed is very similar to that of the raw $\delta^{13}\text{C}$ data.

Fig. S11. Carbon isotope data of the C25, C27 and C29 *n*-alkane ($\delta^{13}\text{C}_{25}$, $\delta^{13}\text{C}_{27}$, $\delta^{13}\text{C}_{29}$) vs. stratigraphic position, Mar 2x section. The shift in isotopic compositions are very similar to bulk $\delta^{13}\text{C}$ data and $\delta^{13}\text{C}_{31}$.

Fig. S12. Deuterium data of *n*-alkane 25, 27, 29, and 31 vs. stratigraphic position, Mar 2x section. A the distinct negative shift can be observed at the onset of the PETM.

Fig. S13. Agglomerative Cluster Analysis for Mar 2x, Rieci Mache and Gonzales. Samples in red belong to PETM interval. Samples in Mar 2x and Gonzales are measured as a well. Samples in Rieci Mache are measured as an outcrop.

Fig. S14. Lithological description of Core P2, San Jacinto-Sinu Belt, northwestern Colombia.

Supplementary Tables

Table S1. Stable carbon isotope ($\delta^{13}\text{C}$) stratigraphy for all studied sections.

Table S2. Biostratigraphic range chart files for Mar 2x (depths are given from the ground-surface).

Table S3. Biostratigraphic range chart files for Rieci Mache (depths are given as measured as an outcrop; greater measure= younger rock).

Table S4. Biostratigraphic range chart files for Gonzales (depths are given from the ground-surface).

Table S5. Analytical data from the U-Pb CA-TIMS zircon analysis of the Rieci Mache Tuff.

Table S6. Analytical data from the U-Pb LA-ICP-MS and CA-TIMS zircon analysis of the Rieci Mache Tuff.

Table S7. Mar 2x carbon isotope composition (C) and deuterium compositions (D) of n-alkanes with 25, 27, 29, and 31 carbon atoms. CPI=Carbon Preference Index.

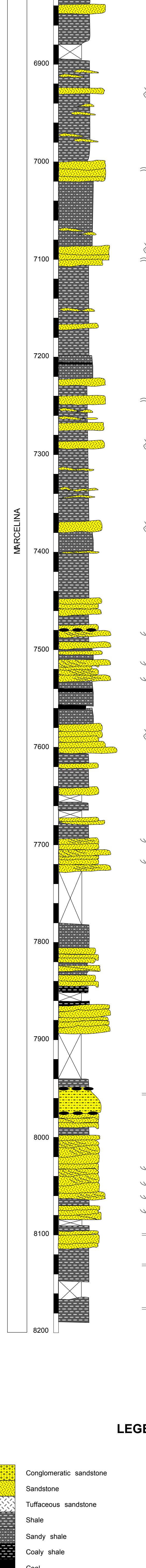
Table S8. Diversity summary results for Mar2x, Rieci Mache and Gonzales sites.
N=number of pollen and spores counted per sample; S=number of species per sample;
SD=Standing Diversity, after range-through method and excluding singletons, numbers
in red indicate samples with edge-effect that were excluded from standing diversity
analysis; SI=Simpson Index (it was not calculated for samples with counts <80).

Table S9. Relative abundances of dry vs wet indicator for Mar2x core. Precipitation
preferences of families identified in this study are derived from Punyasena study (S79) of
Gentry's 144-transect neotropical plant database.

Table S10. P2 core TEX_{86} values, BIT indices and SST estimates following the
calibration of Kim et al. (S85) and Liu et al. (S86).

Figure S1. Lithological description of the Mar 2x Core, Maracaibo, Venezuela

**MAR2X WELL STRATIGRAPHIC COLUMN
PALEOCENE - EOCENE INTERVAL
SCALE 1:2000**



LEGEND

[Yellow square]	Conglomeratic sandstone	[Vertical lines icon]	Planar lamination
[Yellow square]	Sandstone	[≈≈ icon]	Ripple cross-lamination
[Yellow square]	Tuffaceous sandstone	[≡≡ icon]	Trough cross-stratification
[Grey square]	Shale	[≡≡≡≡ icon]	Planar cross-stratification
[Grey square]	Sandy shale	[≈≈≈≈ icon]	Convolute lamination
[Grey square]	Coaly shale	[Leaf icon]	Plant debris
[Black square]	Coal	[Wavy line icon]	Bioturbation
[White square with black X]	Cover		
[Yellow square]	Palynological sample		

Figure S2. Polygonal of the Riecito Mache section, Maracaibo, Venezuela.

SCALE 1:5000

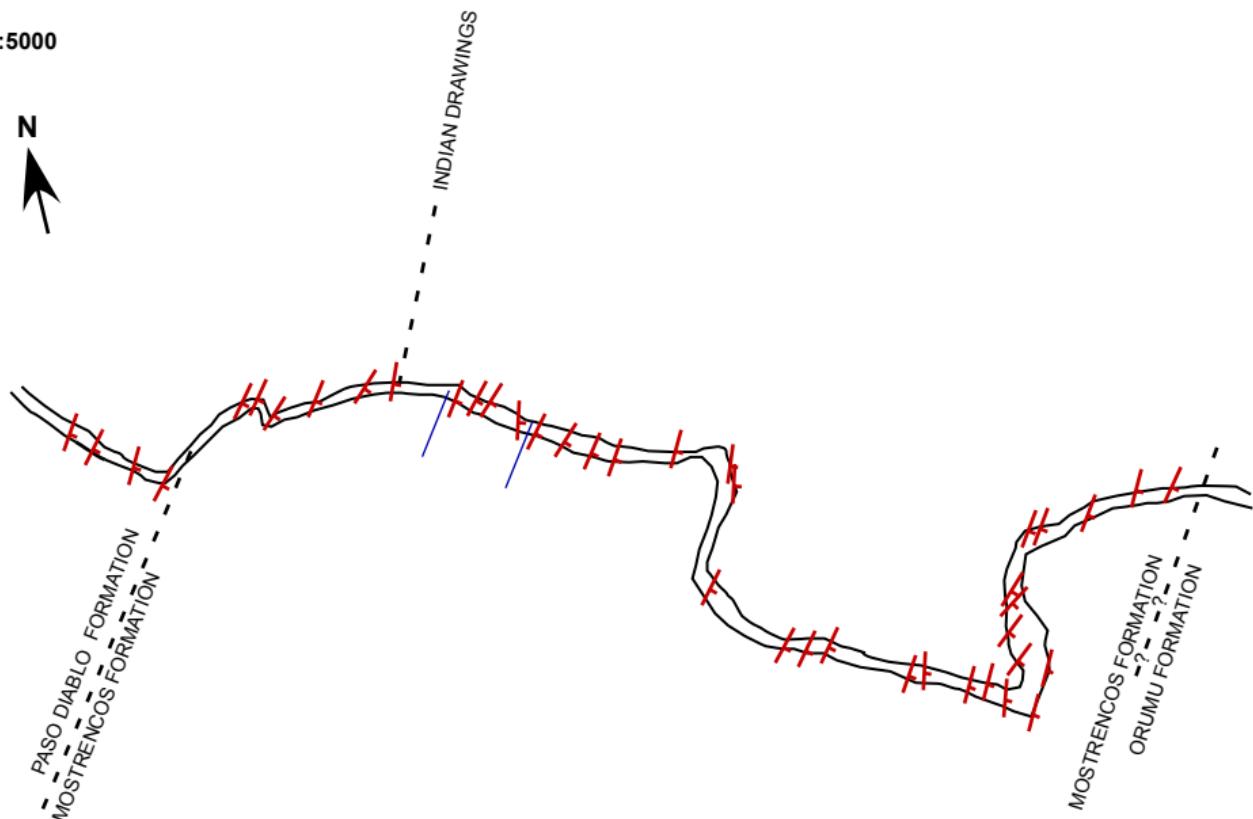
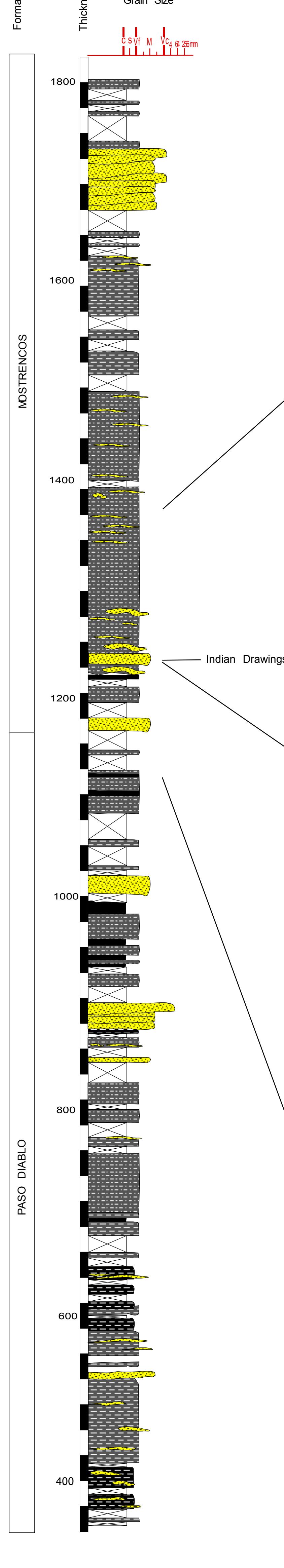


Figure S3. Lithological description of the Rieci Mache, Maracaibo, Venezuela.

**STRATIGRAPHIC COLUMN RIECITO MACHE
PALEOCENE - EOCENE LIMIT**

SECTION BY STEKHOVEN

SCALE 1:2500

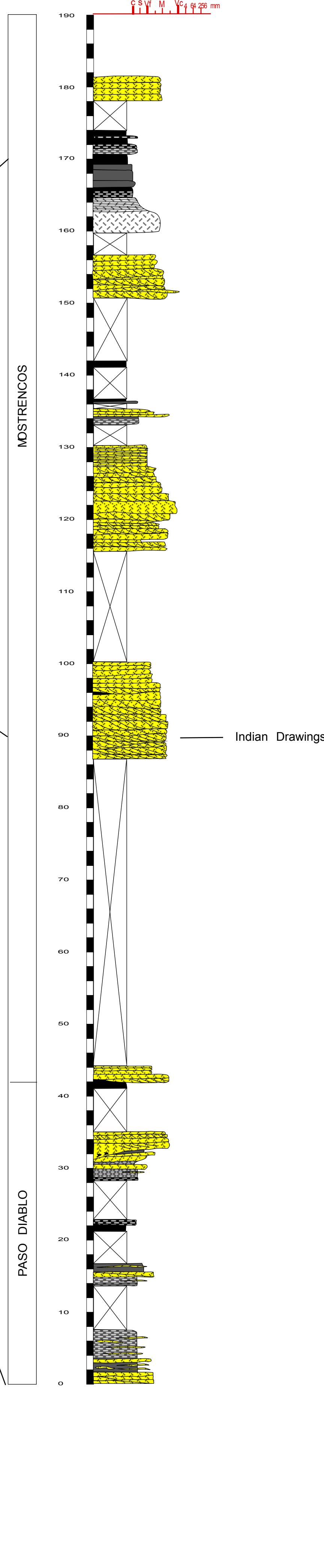


**STRATIGRAPHIC COLUMN RIECITO MACHE
PALEOCENE - EOCENE LIMIT**

ZULIA STATE - VENEZUELA

SECTION BY STRI FIELD TEAM

SCALE 1:200



LEGEND

[Yellow square]	Conglomeratic sandstone	—	Planar lamination
[Yellow dots]	Sandstone	≈	Ripple cross-lamination
[Cross-hatch]	Tuffaceous sandstone	↙	Trough cross-stratification
[Grey]	Shale	//	Planar cross-stratification
[Dotted grey]	Sandy shale	~~	Convolute lamination
[Black]	Coaly shale	⌚	Plant debris
[Coal]	Coal	◎	Bioturbation
[White]	Cover	●	Coal lens

Figure S4. Polygonal of the Gonzales section, Catatumbo, Colombia.

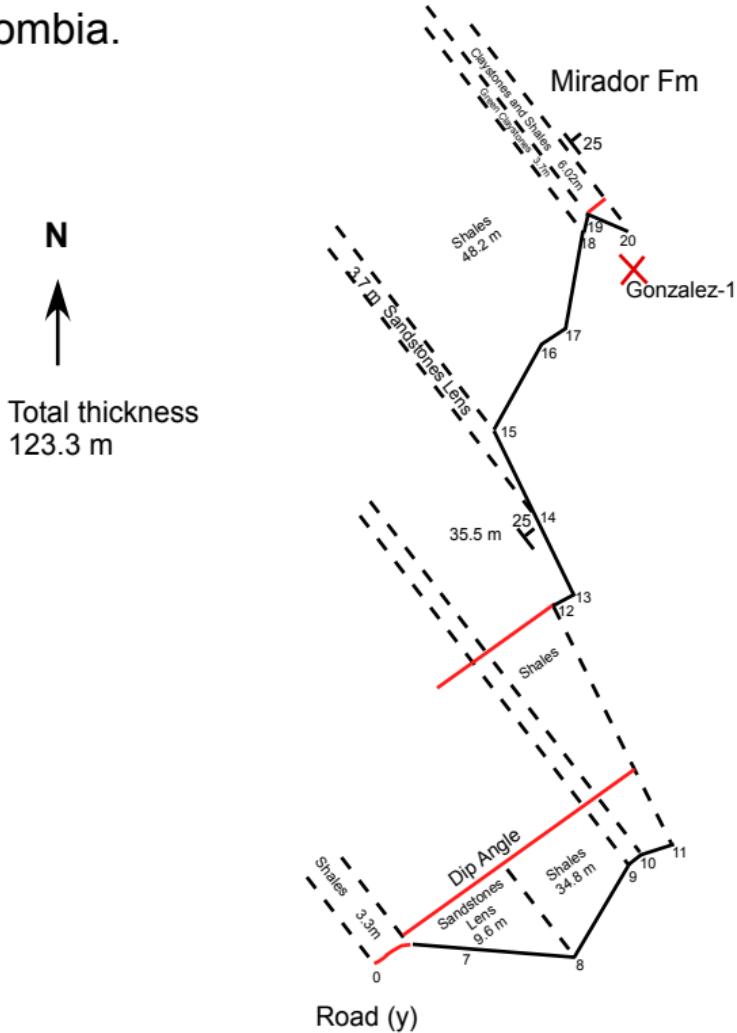
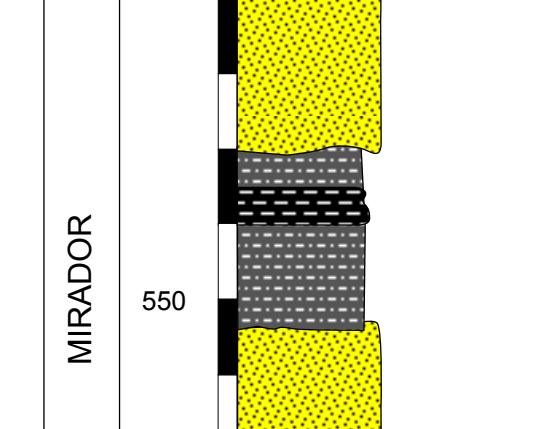


Figure S5. Lithological description of the Gonzales outcrop and Gonzales well, Cataumbo, Colombia.

**STRATIGRAPHIC COLUMN GONZALES-1 WELL
PALEOCENE - EOCENE INTERVAL**

SCALE 1:200

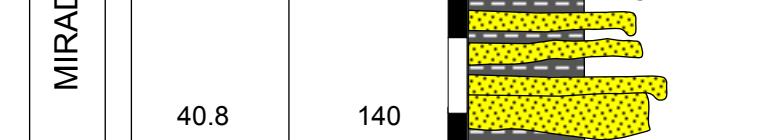


MIRADOR

LOS CUERVOS

**GONZALES OUTCROP STRATIGRAPHIC COLUMN
PALEOCENE - EOCENE INTERVAL**

SCALE 1:200



MIRADOR

LOS CUERVOS

MIRADOR

<p

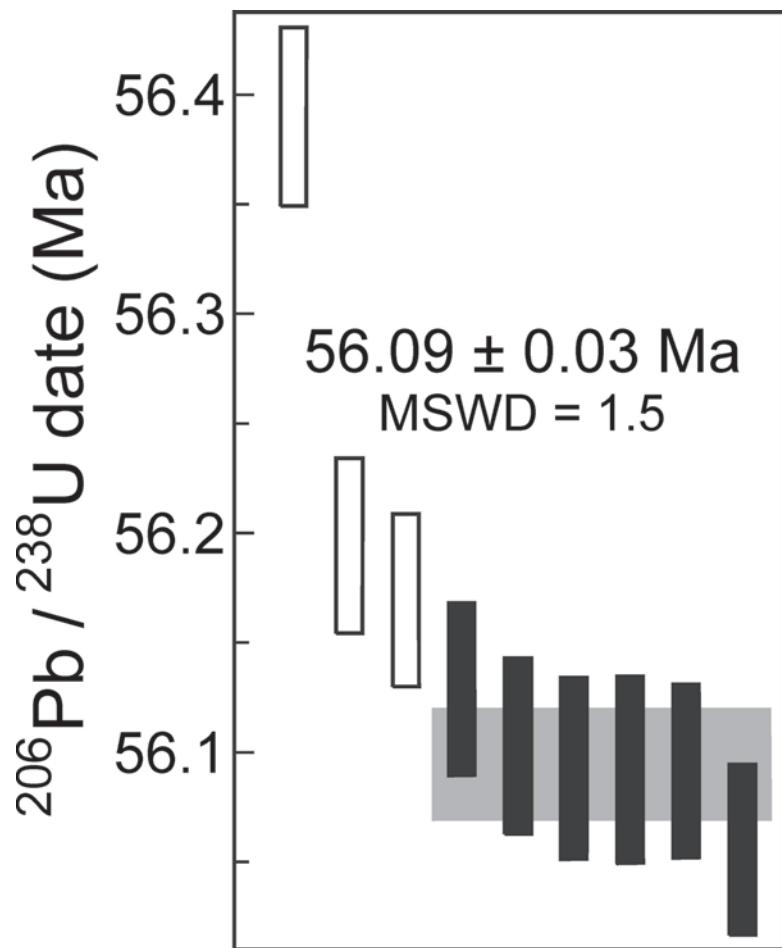


Fig. S6. Plot of the ^{206}Pb - ^{238}U dates from the youngest zircon from the Rieci Mache tuff obtained by the U-Pb CA-TIMS method. Plotted with Isoplot 3.0 (S23). Error bars are 2σ internal errors. Weighted mean date is represented by the grey box behind the error bars. Error bars in white were not used in the weighted mean calculation.

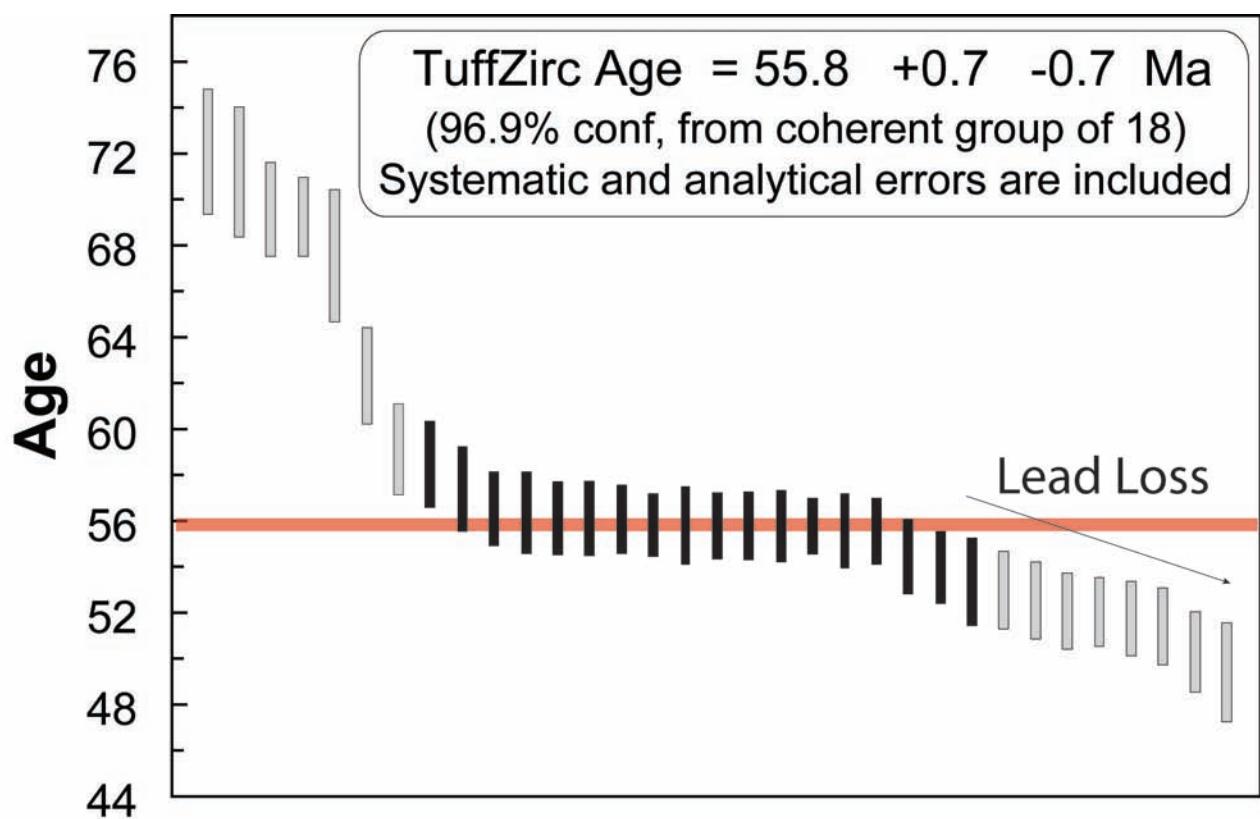


Fig. S7. U-Pb LA-ICP-MS TUFFZIRC age from the Rieciro Mache tuff. Plotted with Isoplot 3.0 (S23).

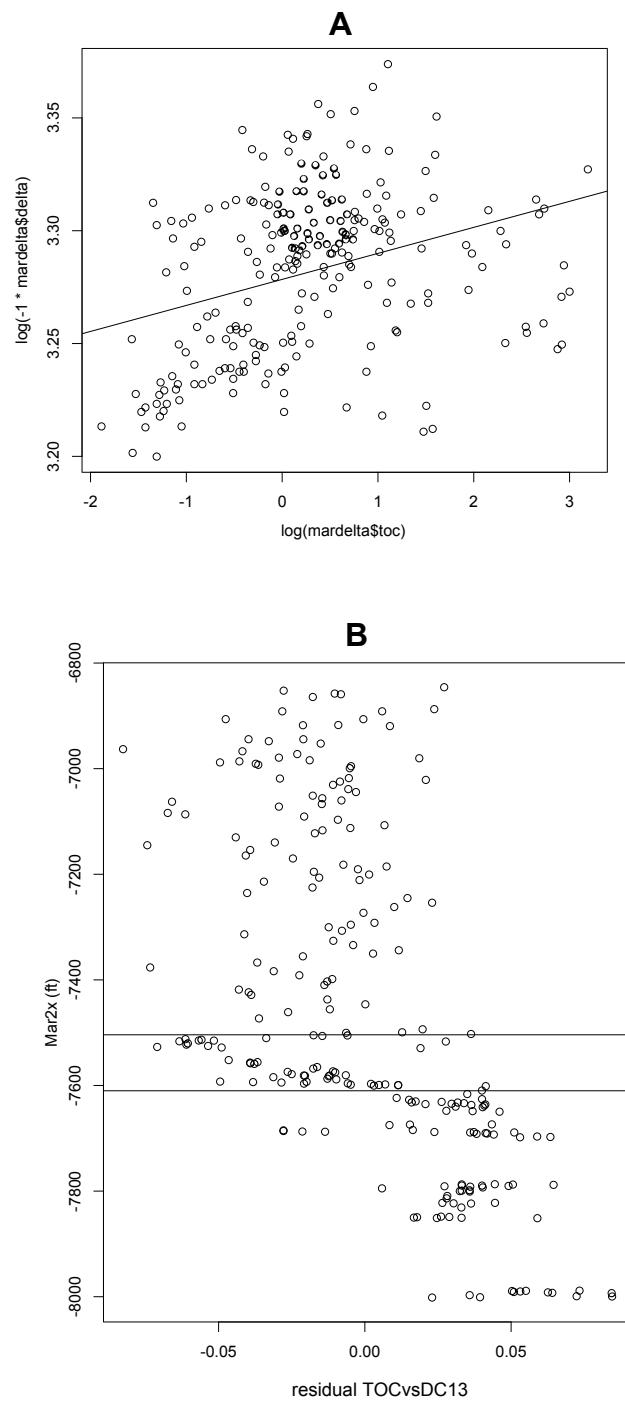


Fig. S8 Carbon isotope data ($\delta^{13}\text{C}$) vs. Stratigraphic position, Mar 2x section following the method by Wing et al (S15). A. Total organic carbon (TOC) vs. carbon isotope data ($\delta^{13}\text{C}$). B. Residual of correlation TOC - $\delta^{13}\text{C}$ vs. the stratigraphic position. Pattern is very similar to raw $\delta^{13}\text{C}$ data.

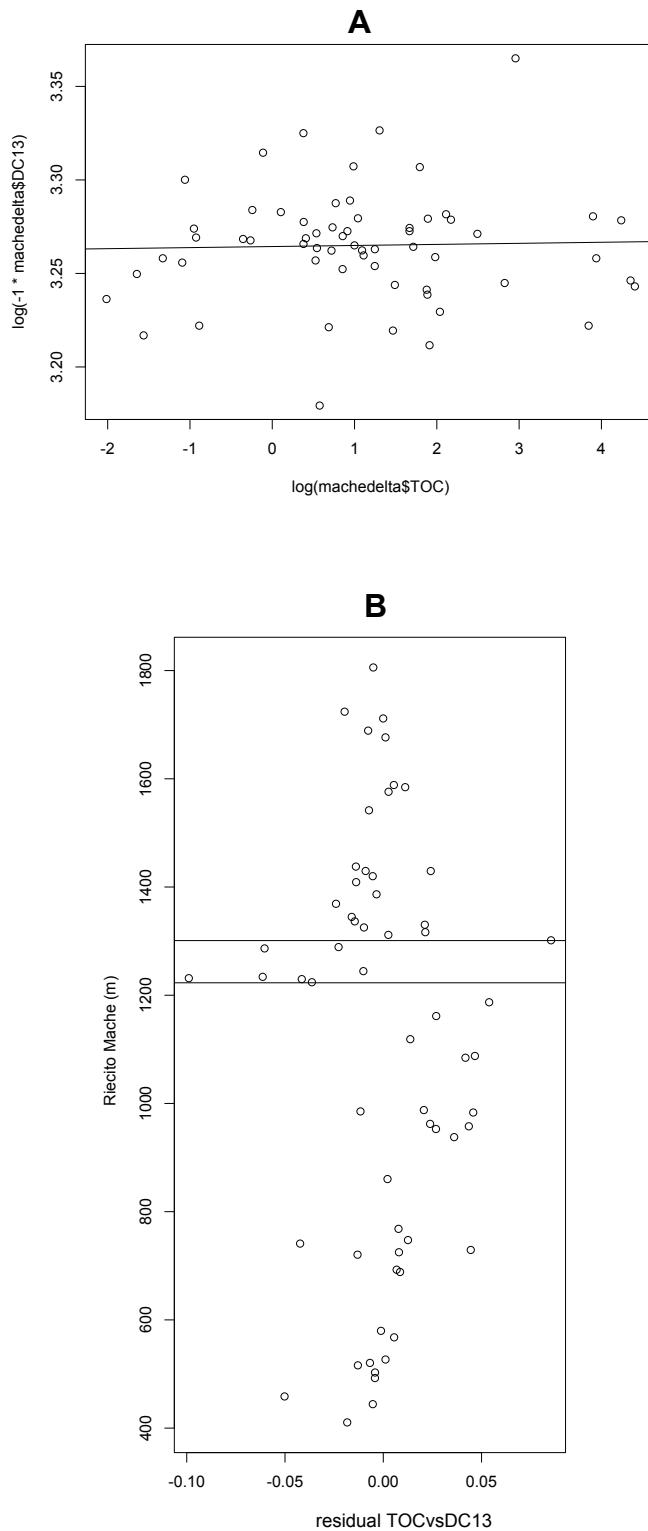


Fig. S9. Carbon isotope data ($\delta^{13}\text{C}$) vs. stratigraphic position, Riecito Mache section following the method by Wing et al (S15). A. Total organic carbon (TOC) vs. carbon isotope data ($\delta^{13}\text{C}$). B. Residual of correlation TOC - $\delta^{13}\text{C}$ vs. the stratigraphic position. Pattern is very similar to raw $\delta^{13}\text{C}$ data.

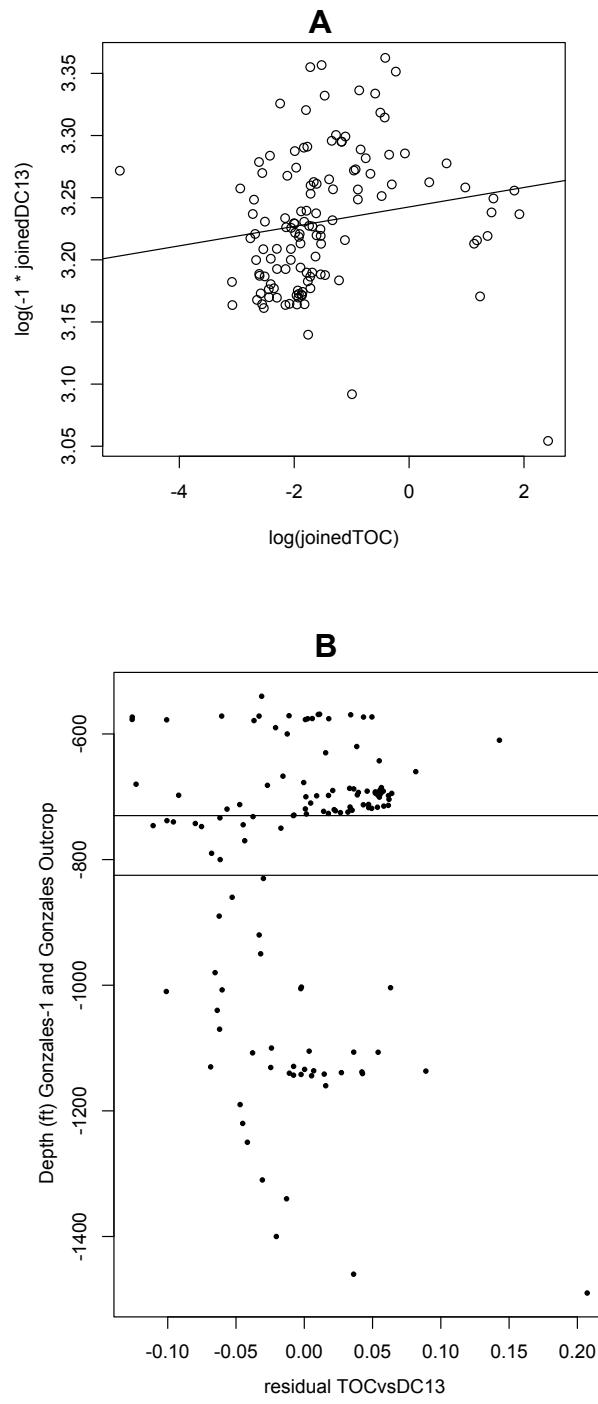


Fig. S10. Carbon isotope data ($\delta^{13}\text{C}$) vs. stratigraphic position, Gonzales section following the method by Wing et al (S15). A. Total organic carbon (TOC) vs. carbon isotope data ($\delta^{13}\text{C}$). B. Residual of correlation TOC - $\delta^{13}\text{C}$ vs. the stratigraphic position. The pattern observed is very similar to that of the raw $\delta^{13}\text{C}$ data.

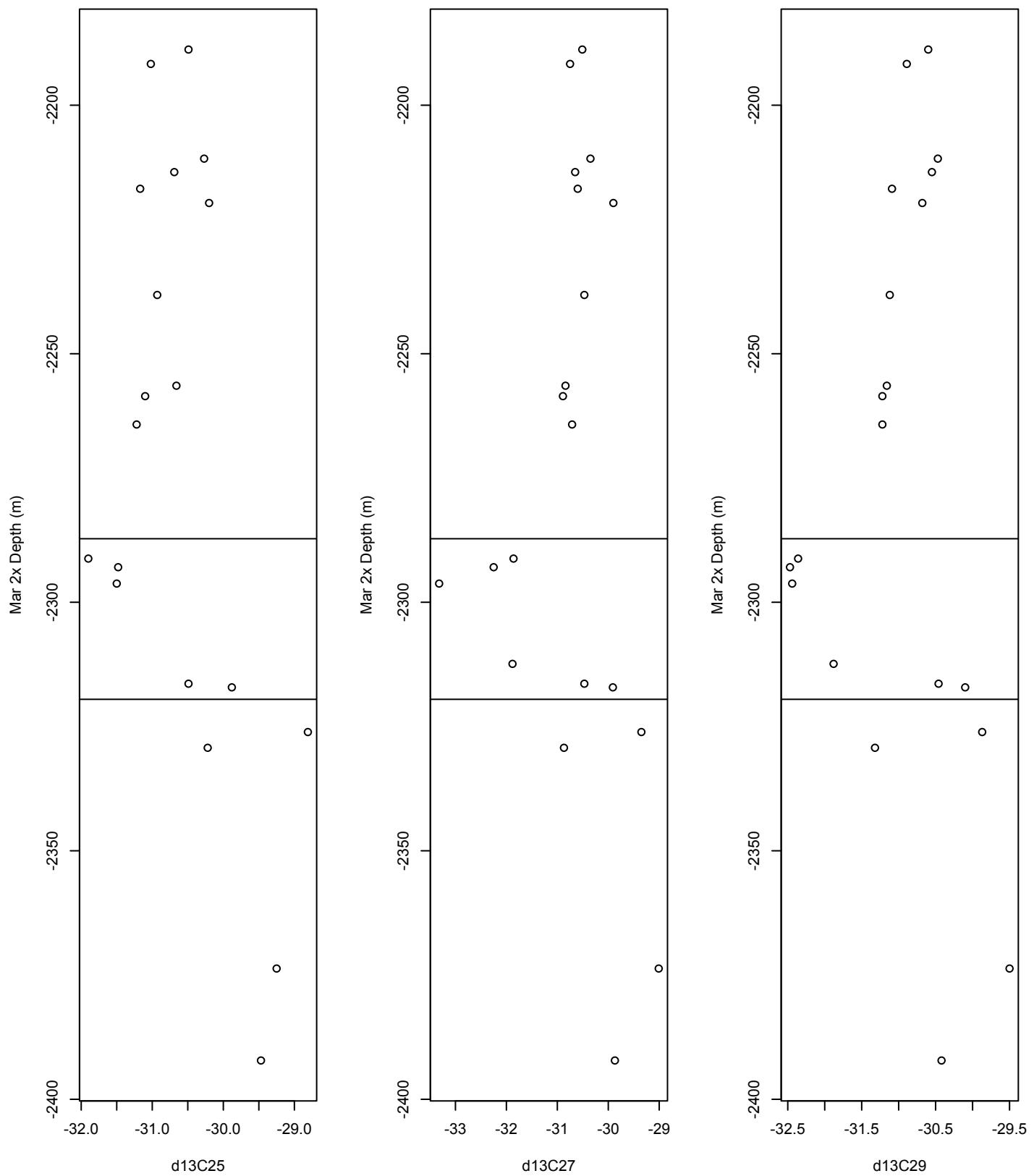


Fig. S11. Carbon isotope data of the C27, C27, and C29 *n*-alkanes ($\delta^{13}\text{C}_{25}$, $\delta^{13}\text{C}_{27}$, $\delta^{13}\text{C}_{29}$) vs. stratigraphic position, Mar 2x section. The shift in isotopic compositions are very similar to bulk $\delta^{13}\text{C}$ data and $\delta^{13}\text{C}_{31}$.

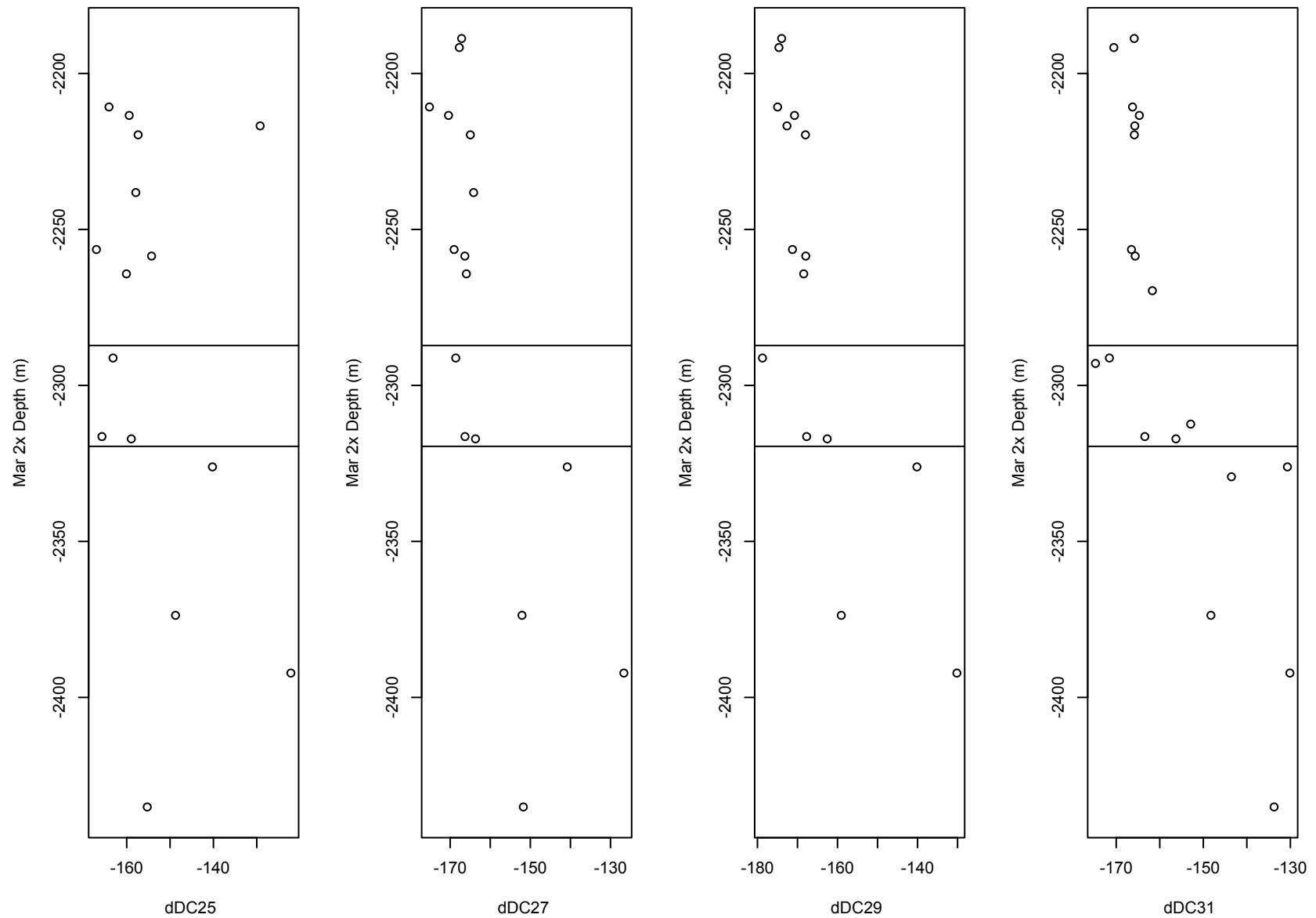


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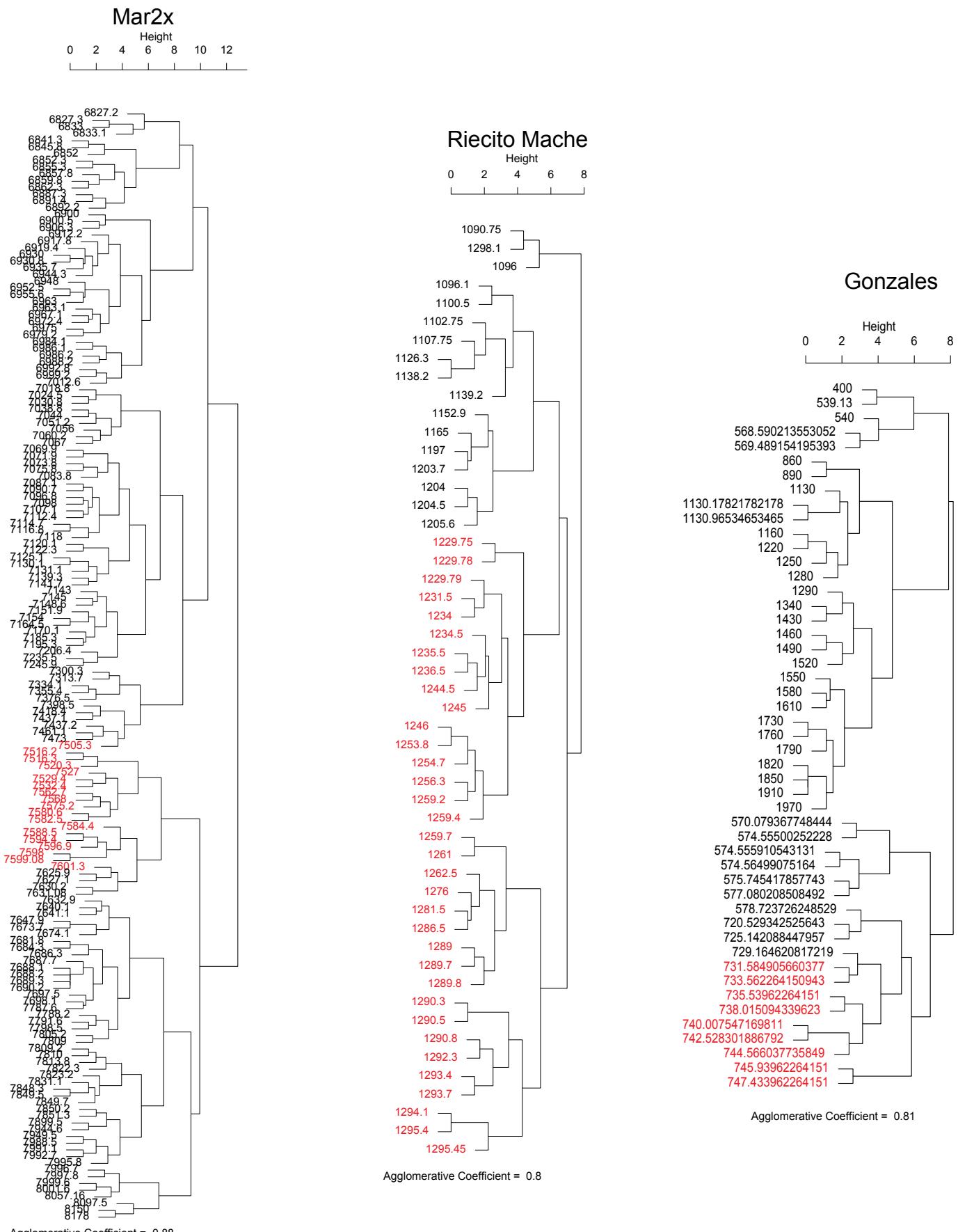


Figure S13. Agglomerative Cluster Analysis for Mar2x, Riequito Mache and Gonzales.
 Samples in red belong to PETM interval. Samples in Mar2x and Gonzales are measured as a well.
 Samples in Riequito Mache are measured as an outcrop.

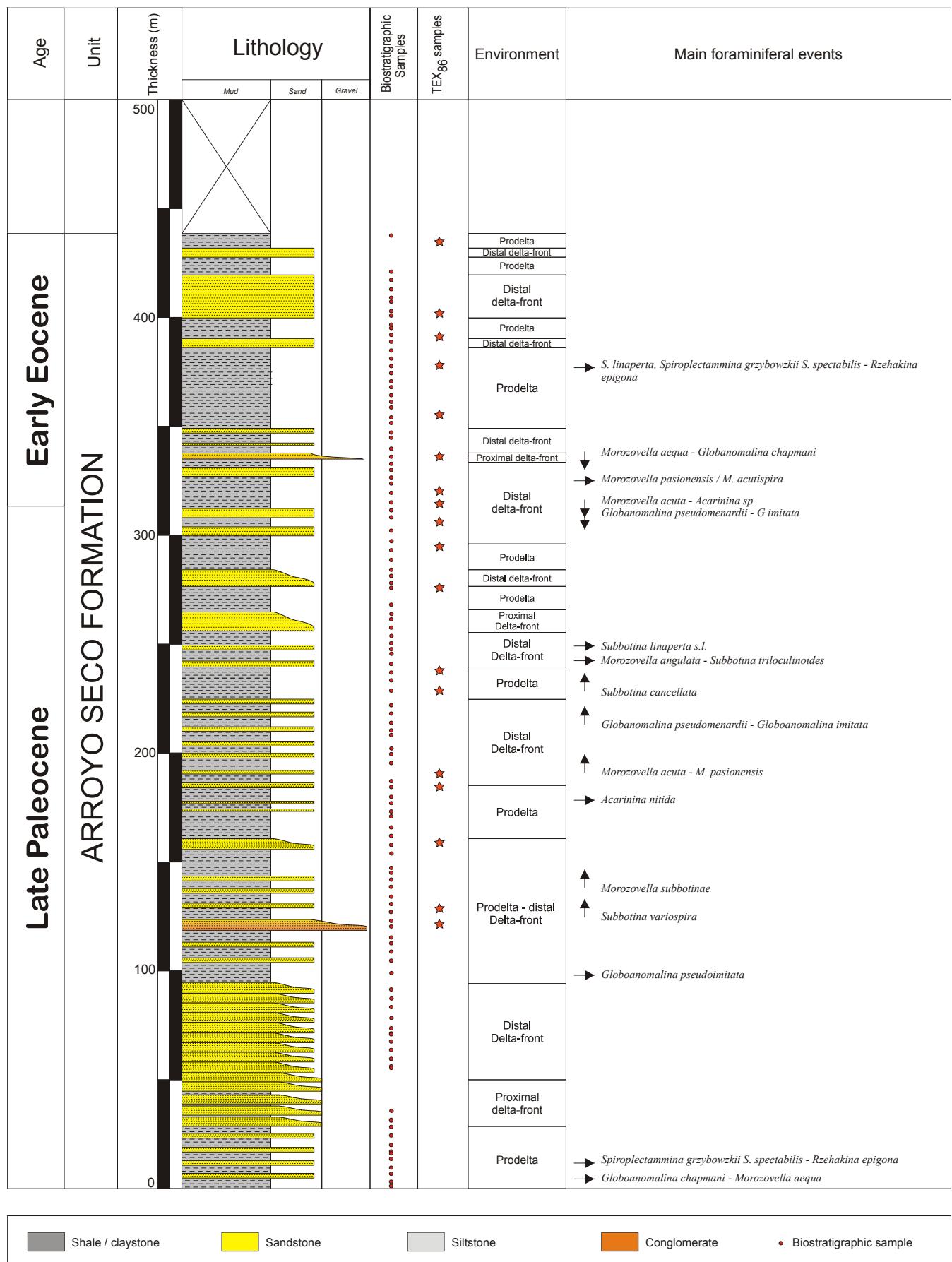


Figure S14. Lithological description of the P2 Core, San Jacinto-Sinu Belt, northwestern Colombia, including key biostratigraphic events, depositional environments and the stratigraphic position of the samples analyzed for biostratigraphy and TEX₈₆

Table S1. Stable carbon isotope stratigraphy for all studied sections.

MAR2x	RIECITO MACHE				GONZALES				
depth (ft)	depth (m)	DC13	TOC	Sample ID	depth (m)	e depth (ft)	e depth (m)	DC13	TOC
6944.3	2116.6	-27.31	1.9662	SS 1361	410.5	-26.65	1.1099	gonzales well-540	164.59
6845.8	2086.6	-26.41	6.9965	SS 1355	444.25	-26.29	0.3963	gonzales well-590	179.83
6852.3	2086.6	-27.56	2.4186	SS 1354	458.5	-27.51	0.894	gonzales well-600	182.88
6851.9	2090.7	-26.83	1.2873	SS 1349	492.5	-26.27	0.7013	gonzales well-610	185.93
6864.3	2092.2	-27.26	2.2205	SS 1348	502.75	-26.28	1.5042	gonzales well-620	188.98
6887.333	2099.3	-25.76	0.6003	SS 1347	516	-26.51	1.465	gonzales well-630	192.02
6891.4	2100.5	-26.56	1.8172	SS 1346	520.5	-26.35	1.7101	gonzales well-660	201.17
6891.4167	2100.5	-27.49	1.8627	SS 1345	526.75	-26.142	1.7211	gonzales well-680	207.26
6906.25	2105	-28.11	2.4065	SS 1342	567.75	-26	0.2644	gonzales well-690	210.31
6906.3	2105	-26.86	2.7573	SS 1340	579.75	-26.2	1.4612	gonzales well-700	213.36
6917.4	2108.4	-26.96	1.8102	SS 2172	688.5	-26	51.291	gonzales well-710	216.41
6917.75	2108.5	-27.34	2.1359	SS 1331	692.75	-26.016	7.2465	gonzales well-720	219.46
6919.4167	2109	-26.37	1.2286	SS 1328	720.5	-26.542	8.773	gonzales well-730	222.5
6944.25	2116.6	-27.79	1.754	SS 1327	725	-25.94	0.3349	gonzales well-750	228.6
6948	2117.8	-27.55	1.5081	SS 2168	729.25	-25.08	46.751	gonzales well-770	234.7
6952.5	2119.1	-26.9	0.8861	SS 1324	741	-27.31	2.6828	gonzales well-790	240.79
6963.0833	2122.3	-26.19	3.016	SS 1322	747.5	-25.8	2.3506	gonzales well-800	243.84
6967.0833	2123.6	-27.74	1.2525	SS 1319	768.25	-25.97	1.6924	gonzales well-830	252.98
6972.4167	2125.2	-27.02	0.6505	SS 1313	860.25	-26.126	3.4788	gonzales well-860	262.13
6979.1667	2127.3	-27.31	0.9525	SS 1306	937.75	-25.267	7.7	gonzales well-890	271.27
6980.3	2127.6	-26.68	8.0803	SS 1305	952.75	-25.499	6.589	gonzales well-920	280.42
6984.1	2128.8	-27.23	1.8389	SS 1304	957.75	-25.059	1.987	gonzales well-950	289.56
6986.1667	2129.4	-27.87	1.7163	SS 2161	962.25	-25.612	81.981	gonzales well-980	298.7
6988.1667	2130	-28.02	1.5393	SS 1301	983.25	-25.014	4.334	gonzales well-1011	307.85
6990.7	2130.8	-27.59	1.1616	SS 2159	985.25	-26.533	69.472	gonzales well-1041	316.99
6992.75	2131.4	-27.59	1.2517	SS 2158	987.75	-25.693	77.802	gonzales well-1071	326.14
6999.1667	2133.3	-26.72	1.1716	SS 1296	1084.5	-25.08	0.4116	gonzales well-1101	335.28
7017.9167	2139.1	-26.84	1.6819	SS 1295	1087.75	-24.95	0.2094	gonzales well-1131	344.42
7018.75	2139.3	-27.48	1.6819	SS 1294	1090.75	-24.433		gonzales well-1161	353.57
7021.3	2140.1	-26.13	1.6123	SS 1293	1096	-25.09		gonzales well-1191	362.71
7024.5	2141.1	-26.81	1.1725	SS 2153	1125	-24.732		gonzales well-1221	371.86
7030.6667	2142.9	-26.88	1.1968	SS 1291	1107.75	-24.893		gonzales well-1251	381
7038.75	2145.4	-26.84	1.6537	SS 2152	1118	-24.936		gonzales well-1311	399.29
7044.3333	2147.1	-26.65	1.1209	SS 1290	1118.75	-25.783	0.1929	gonzales well-1341	408.43
7051.1667	2149.2	-27.05	1.1339	SS 1289	1161.5	-25.44	0.1334	gonzales well-1401	426.72
7056	2150.7	-27.01	1.3225	SS 1288	1187	-24.819	6.7633	gonzales well-1461	445.01
7060.1667	2151.9	-26.77	1.0761	SS 1287	1197	-25.58		gonzales well-1491	454.15
7062.6667	2152.7	-28.59	2.128	SS 1283	1224	-27.115	0.3461	gonzales outcrop	0.17
7067	2154	-27.05	1.4827	SS 1284	1229.75	-27.299	6.0082	gonzales outcrop	0.37
7071.9167	2155.5	-27.33	1.0152	SS 2150	1231.5	-28.934	19.226	gonzales outcrop	0.6
7083.8333	2159.2	-28.35	1.6599	SS 1282	1234	-27.84	3.6843	gonzales outcrop	0.72
7086.5	2160	-28.3	1.3024	SS 1280	1244.5	-26.416	0.386	gonzales outcrop	0.88
7090.6667	2161.2	-27.14	1.1724	SS 1277	1286.5	-27.799	1.46	gonzales outcrop	1
7088.17	2343.4	-25.92	3.32	SS 1269	1336.5	-26.562	2.8332	gonzales outcrop	1.19
7088.92	2343.6	-25.47	2.41	SS 1267	1344.75	-26.619	8.2654	gonzales outcrop	1.41
7089.17	2343.6	-25.38	0.48	SS 1264	1369	-26.815	2.5721	gonzales outcrop	1.69
7087.17	2343	-27.21	1.39	SS 1272	1316.5	-25.632	4.4364	gonzales outcrop	1.78
7087.67	2343.2	-27.35	4.25	SS 1271	1325	-26.434	2.08	gonzales outcrop	2.24
7088.08	2343.3	-26.37	4.59	SS 1270	1330.25	-26.659	16.85	gonzales outcrop	2.85
7088.17	2343.4	-25.92	3.32	SS 1269	1336.5	-26.562	2.8332	gonzales outcrop	3.2
7088.92	2343.6	-25.47	2.41	SS 1267	1344.75	-26.619	8.2654	gonzales outcrop	7.55
7089.17	2343.6	-25.38	0.48	SS 1264	1369	-26.815	2.5721	gonzales outcrop	7.75
7090.17	2344	-25.11	0.3	SS 1261	1386.5	-26.5	0.7676	gonzales outcrop	7.79
7090.92	2344.2	-25.02	0.23	SS 1260	1409	-26.556	6.6273	gonzales outcrop	8.1
7091.58	2344.4	-25.39	0.6	SS 1258	1420	-26.342	12.095	gonzales outcrop	27.8
7092.92	2344.8	-25.03	0.29	SS 2140	1429.5	-25.567	6.5329	gonzales outcrop	28.2
7096.67	2345.9	-24.57	0.21	SS 1256	1429.6	-26.425	5.3045	gonzales outcrop	28.5
7097.5	2346.2	-24.53	0.27	SS 2139	1437.75	-26.59	49.288	gonzales outcrop	28.7
7098.08	2346.4	-24.86	0.35	SS 1243	1541.75	-26.38	5.3013	gonzales outcrop	80.3
7098.08	2346.4	-25.37	4.59	SS 1270	1330.25	-26.659	16.85	gonzales outcrop	8.6
7098.17	2343.4	-25.92	3.32	SS 1269	1336.5	-26.562	2.8332	gonzales outcrop	8.7
7098.92	2343.6	-25.47	2.41	SS 1267	1344.75	-26.619	8.2654	gonzales outcrop	7.55
7098.92	2343.6	-25.38	0.48	SS 1264	1369	-26.815	2.5721	gonzales outcrop	7.75
7100.25	2345.9	-25.11	0.3	SS 1261	1386.5	-26.5	0.7676	gonzales outcrop	7.79
7100.92	2346.2	-25.02	0.23	SS 1260	1409	-26.556	6.6273	gonzales outcrop	8.1
7101.58	2344.4	-25.39	0.6	SS 1258	1420	-26.342	12.095	gonzales outcrop	27.8
7102.92	2344.8	-25.03	0.29	SS 2140	1429.5	-25.567	6.5329	gonzales outcrop	28.2
7106.67	2345.9	-24.57	0.21	SS 1256	1429.6	-26.425	5.3045	gonzales outcrop	28.5
7107.5	2346.2	-24.53	0.27	SS 2139	1437.75	-26.59	49.288	gonzales outcrop	28.7
7109.25	2347.1	-25.27	0.33	SS 1224	1724	-26.68	0.7849	gonzales outcrop	80.3
7109.92	2347.5	-25.07	0.24	SS 1228	1805.75	-26.31	2.3542	gonzales outcrop	80.88
7110.25	2347.7	-25.47	0.64					gonzales outcrop	84.1
7110.92	2347.7	-25.33	0.55					gonzales outcrop	84.7
7110.92	2347.7	-25.33	0.4					gonzales outcrop	85
7110.92	2347.7	-25.33	0.3					gonzales outcrop	85.4
7110.92	2347.7	-25.33	0.2					gonzales outcrop	85.7
7110.92	2347.7	-25.33	0.1					gonzales outcrop	86.1
7110.92	2347.7	-25.33	0.0					gonzales outcrop	86.4
7110.92	2347.7	-25.33	-0.1					gonzales outcrop	86.7
7110.92	2347.7	-25.33	-0.2					gonzales outcrop	87

Table S5. Analytical data from the U-Pb CA-TIMS zircon analysis of the Rieciro Mache Tuff.

Sample	Radiogenic Isotope Ratios													Isotopic Ages						
	<u>Th</u>	$^{206}\text{Pb}^*$	mol %	<u>Pb*</u>	<u>Pb_c</u>	$\frac{^{206}\text{Pb}}{^{204}\text{Pb}}$	$\frac{^{208}\text{Pb}}{^{206}\text{Pb}}$	$\frac{^{207}\text{Pb}}{^{206}\text{Pb}}$	% err	$\frac{^{207}\text{Pb}}{^{235}\text{U}}$	$\frac{^{206}\text{Pb}}{^{238}\text{U}}$	corr.	$\frac{^{207}\text{Pb}}{^{206}\text{Pb}}$	\pm	$\frac{^{207}\text{Pb}}{^{235}\text{U}}$	\pm	$\frac{^{206}\text{Pb}}{^{238}\text{U}}$	\pm		
	(a)	(b)	(c)	(c)	(c)	(d)	(e)	(e)	(f)	(e)	(f)	(f)	(g)	(f)	(g)	(f)	(g)	(f)		
z1	0.494	0.8933	99.63%	81	0.27	4989	0.159	0.047122	0.129	0.056884	0.196	0.008755	0.065	0.776	55.40	3.08	56.18	0.11	56.19	0.04
z2*	0.426	0.7152	99.40%	50	0.35	3122	0.137	0.047072	0.212	0.056731	0.258	0.008741	0.072	0.721	52.90	5.05	56.03	0.14	56.10	0.04
z3*	0.469	0.4306	99.26%	41	0.26	2528	0.150	0.047017	0.300	0.056654	0.352	0.008739	0.074	0.761	50.09	7.15	55.96	0.19	56.09	0.04
z5*	0.420	1.4577	99.74%	112	0.32	7064	0.134	0.047066	0.130	0.056751	0.183	0.008745	0.070	0.840	52.60	3.10	56.05	0.10	56.13	0.04
z6	0.335	2.3849	99.79%	141	0.41	8957	0.116	0.051418	0.208	0.074659	0.270	0.010531	0.139	0.651	259.60	4.78	73.11	0.19	67.53	0.09
z7*	0.440	2.5740	99.74%	116	0.54	7260	0.141	0.047141	0.113	0.056803	0.165	0.008739	0.071	0.839	56.40	2.69	56.10	0.09	56.09	0.04
z8	0.454	0.8794	99.60%	75	0.29	4697	0.145	0.047130	0.166	0.056869	0.214	0.008751	0.071	0.772	55.83	3.95	56.16	0.12	56.17	0.04
z9*	0.371	1.1111	99.63%	78	0.34	4999	0.119	0.047217	0.144	0.056857	0.194	0.008733	0.070	0.807	60.22	3.42	56.15	0.11	56.06	0.04
z11*	0.408	0.7796	99.40%	49	0.39	3087	0.131	0.047155	0.224	0.056820	0.270	0.008739	0.077	0.695	57.08	5.34	56.11	0.15	56.09	0.04
z12	0.476	0.8707	99.48%	58	0.37	3574	0.153	0.047245	0.176	0.057232	0.226	0.008786	0.073	0.764	61.61	4.20	56.51	0.12	56.39	0.04

(a) z1, z2, etc. are labels for analyses composed of single zircon grains that were annealed and chemically abraded (S32). Fraction labels with * denote analyses used in the weighted mean calculation.

(b) Model Th/U ratio calculated from radiogenic $^{208}\text{Pb}/^{206}\text{Pb}$ ratio and $^{207}\text{Pb}/^{235}\text{U}$ date.

(c) Pb* and Pbc are radiogenic and common Pb, respectively. mol % $^{206}\text{Pb}^*$ is with respect to radiogenic and blank Pb.

(d) Measured ratio corrected for spike and fractionation only. Fractionation correction is 0.15 ± 0.03 (1 sigma) %/amu (atomic mass unit) for single-collector Daly analyses, based on analysis of EARTHTIME 202Pb-205Pb tracer solution.

(e) Corrected for fractionation, spike, common Pb, and initial disequilibrium in $^{230}\text{Th}/^{238}\text{U}$. Common Pb is assigned to procedural blank with composition of $^{206}\text{Pb}/^{204}\text{Pb} = 18.60 \pm 0.80\%$; $^{207}\text{Pb}/^{204}\text{Pb} = 15.69 \pm 0.32\%$; $^{208}\text{Pb}/^{204}\text{Pb} = 38.51 \pm 0.74\%$ (1 sigma). $^{206}\text{Pb}/^{238}\text{U}$ and $^{207}\text{Pb}/^{206}\text{Pb}$ ratios corrected for initial disequilibrium in $^{230}\text{Th}/^{238}\text{U}$ using Th/U [magma] = 3.

(f) Errors are 2 sigma, propagated using algorithms of Schmitz and Schoene (S35) and Crowley et al. (S36).

(g) Calculations based on the decay constants of Jaffey et al. (S24). $^{206}\text{Pb}/^{238}\text{U}$ and $^{207}\text{Pb}/^{206}\text{Pb}$ dates corrected for initial disequilibrium in $^{230}\text{Th}/^{238}\text{U}$ using Th/U [magma] = 3.

Table S6. Analytical data from the U-Pb LA-ICP-MS and CA-TIMS zircon analysis of the Rieciro Mache Tuff.

sample	238U	1 sigma	207Pb	1 sigma	206/238	1 sigma	207/206	1 sigma	Best age	1 sigma
name	206Pb	% error	206Pb	% error	age	abs err	age	abs err	abs err Ma	
B-110_50	129.9825	2.20%	0.0477	3.28%	49.4	1.1	252.3	69.7	49.4	1.1
B-110_49	127.7320	1.76%	0.0455	2.09%	50.3	0.9	0.0	24.1	50.3	0.9
B-110_48	124.9247	1.65%	0.0476	1.83%	51.4	0.8	43.9	41.7	51.4	0.8
B-110_47	124.0451	1.57%	0.0468	1.58%	51.8	0.8	39.5	39.9	51.8	0.8
B-110_46	123.3919	1.44%	0.0466	1.52%	52.0	0.7	28.9	38.5	52.0	0.7
B-110_45	123.3185	1.61%	0.0454	1.33%	52.1	0.8	0.0	0.0	52.1	0.8
B-110_44	122.2317	1.62%	0.0485	1.67%	52.5	0.8	158.1	38.5	52.5	0.8
B-110_43	121.1531	1.61%	0.0448	1.51%	53.0	0.8	0.0	0.0	53.0	0.8
B-110_42 *	120.3530	1.81%	0.0475	2.27%	53.3	1.0	66.9	54.9	53.3	1.0
B-110_41 *	118.9386	1.45%	0.0474	1.55%	54.0	0.8	93.1	38.4	54.0	0.8
B-110_40 *	117.9010	1.51%	0.0477	1.40%	54.4	0.8	83.4	34.4	54.4	0.8
B-110_39 *	115.5159	1.31%	0.0468	0.90%	55.6	0.7	73.2	27.8	55.6	0.7
B-110_38 *	115.4896	1.48%	0.0484	1.79%	55.6	0.8	146.8	43.8	55.6	0.8
B-110_37 *	115.0622	1.11%	0.0457	0.86%	55.8	0.6	0.0	15.9	55.8	0.6
B-110_36 *	115.0535	1.42%	0.0480	1.42%	55.8	0.8	99.0	35.9	55.8	0.8
B-110_35 *	115.0333	1.32%	0.0467	1.27%	55.8	0.7	39.4	33.4	55.8	0.7
B-110_34 *	115.0225	1.30%	0.0477	1.29%	55.8	0.7	167.1	32.5	55.8	0.7
B-110_33 *	114.9871	1.53%	0.0472	1.67%	55.8	0.9	73.8	43.2	55.8	0.9
B-110_32 *	114.9507	1.24%	0.0463	1.30%	55.8	0.7	63.0	33.7	55.8	0.7
B-110_31 *	114.4626	1.33%	0.0469	1.08%	56.1	0.7	61.0	31.3	56.1	0.7
B-110_30 *	114.3735	1.47%	0.0462	1.14%	56.1	0.8	17.8	30.9	56.1	0.8
B-110_29 *	114.3663	1.42%	0.0469	1.15%	56.1	0.8	87.0	31.2	56.1	0.8
B-110_28 *	113.8808	1.59%	0.0467	1.26%	56.4	0.9	31.5	31.7	56.4	0.9
B-110_27 *	113.5241	1.44%	0.0469	1.41%	56.5	0.8	99.6	34.3	56.5	0.8
B-110_26 *	111.8125	1.62%	0.0465	1.05%	57.4	0.9	23.9	28.3	57.4	0.9
B-110_25 *	109.7222	1.63%	0.0463	2.20%	58.5	0.9	162.2	59.3	58.5	0.9
B-110_24	108.4996	1.69%	0.0513	1.70%	59.1	1.0	393.1	44.7	59.1	1.0
B-110_23	102.8870	1.71%	0.0486	1.73%	62.4	1.1	128.2	42.2	62.4	1.1
B-110_22	94.8850	2.15%	0.0531	2.61%	67.6	1.4	332.8	59.7	67.6	1.4
B-110_21	92.5295	1.26%	0.0477	1.21%	69.3	0.9	79.0	32.1	69.3	0.9
B-110_20	92.1185	1.49%	0.0455	1.47%	69.6	1.0	128.9	47.5	69.6	1.0
B-110_19	89.9886	2.01%	0.0483	2.53%	71.2	1.4	177.0	61.2	71.2	1.4
B-110_18	88.8842	1.92%	0.0445	2.02%	72.1	1.4	0.0	0.0	72.1	1.4
B-110_17	68.0999	1.52%	0.0492	1.81%	94.0	1.4	158.2	44.0	94.0	1.4
B-110_16	66.7055	1.30%	0.0474	1.15%	95.9	1.2	114.4	35.2	95.9	1.2
B-110_15	31.4134	1.05%	0.0502	0.63%	202.0	2.1	205.1	17.8	202.0	2.1
B-110_14	26.3220	1.12%	0.0505	0.76%	240.4	2.6	220.0	21.8	240.4	2.6
B-110_13	25.8561	3.29%	0.0520	3.32%	244.6	7.9	286.2	74.9	244.6	7.9
B-110_12	23.2769	1.30%	0.0518	0.66%	271.2	3.4	275.6	18.5	271.2	3.4
B-110_11	22.7715	1.21%	0.0592	1.03%	277.1	3.3	643.9	24.0	277.1	3.3
B-110_10	22.3345	1.63%	0.0520	1.63%	282.4	4.5	278.8	38.9	282.4	4.5
B-110_9	21.7107	1.18%	0.0526	0.67%	290.3	3.3	313.6	19.2	290.3	3.3
B-110_8	13.8995	1.03%	0.0560	0.54%	447.9	4.5	453.1	15.4	447.9	4.5
B-110_7	7.1388	1.88%	0.0740	0.80%	845.1	14.9	1030.1	19.4	845.1	14.9
B-110_6	7.0245	1.97%	0.0690	0.97%	858.0	15.8	900.8	21.9	858.0	15.8
B-110_5	5.9649	1.63%	0.0726	1.09%	999.1	15.0	1028.7	26.2	1028.7	26.2
B-110_4	5.1921	1.11%	0.0787	0.55%	1135.5	11.5	1166.6	15.7	1166.6	15.7
B-110_3	5.0852	1.14%	0.0788	0.59%	1157.3	12.1	1166.7	14.7	1166.7	14.7
B-110_2	5.3680	2.21%	0.0815	1.04%	1101.2	22.3	1256.4	24.3	1256.4	24.3

BOLD * = Analyzed accepted in the TUFFZIRC age calculation

U/Pb and 206Pb/207Pb fractionation is calibrated relative to fragments of a large Peixe zircon of 564 ± 4 Ma (2-sigma).

All uncertainties are reported at the 1-sigma level, and include only measurement errors. Systematic errors would increase age uncertainties by 1-2%.

Best age = $^{206}\text{Pb} / ^{238}\text{U}$ for ages <1000 Ma and $^{206}\text{Pb} / ^{207}\text{Pb}$ for ages >1000 Ma

Table S7. Mar2x DC13 isotopic composition (C) and Deuterium compositions (D) of n-alkanes with 25, 27, 29, and 31 carbon atoms. CPI=Carbon Preference Index

Depth (ft)	Depth (m)	CPI (ΣOdd(C25-C33)/ΣEven(C26-C34))	Average Chain Length	Isotopic Compositions (δ)							
				δ ¹³ C ₂₅	δ ¹³ C ₂₇	δ ¹³ C ₂₉	δ ¹³ C ₃₁	δD _{n-C17}	δD _{n-C25}	δD _{n-C27}	δD _{n-C29}
7181.10	2188.80	1.47	29.97	-30.49	-30.51	-30.60	-31.62		-167.15	-173.96	-165.87
7190.60	2191.69	1.50	30.05	-31.02	-30.75	-30.89	-31.72		-167.72	-174.63	-170.55
7253.11	2210.75	1.58	30.23	-30.27	-30.35	-30.47	-31.44	-164.08	-175.18	-174.96	-166.25
7262.00	2213.46	1.66	29.81	-30.69	-30.65	-30.55	-31.31	-159.42	-170.42	-170.74	-164.69
7273.00	2216.81	1.61	29.91	-31.17	-30.60	-31.09	-32.32	-129.23		-172.63	-165.74
7282.40	2219.68	1.49	29.85	-30.20	-29.90	-30.68	-31.44	-157.37	-164.97	-168.00	-165.84
7343.10	2238.18	1.61	29.81	-30.93	-30.47	-31.12	-32.30	-157.90	-164.16		
7403.00	2256.43	1.56	29.70	-30.66	-30.84	-31.16	-32.03	-166.98	-169.03	-171.25	-166.50
7409.90	2258.54	1.53	29.46	-31.10	-30.89	-31.22	-32.18	-154.26	-166.34	-167.93	-165.64
7428.60	2264.24	1.44	29.43	-31.22	-30.71	-31.22	-32.19	-160.06	-165.96	-168.44	
7446.30	2269.63										-161.70
7517.11	2291.22	1.58	30.00	-31.90	-31.86	-32.36	-33.38	-163.15	-168.62	-178.75	-171.57
7522.80	2292.95	1.56	29.79	-31.48	-32.25	-32.47	-33.53				-174.77
7533.60	2296.24	1.29	29.35	-31.50	-33.32	-32.44	-32.95				
7586.60	2312.40	1.42	29.60		-31.88	-31.88	-33.10	-165.9			-152.90
7599.70	2316.39	1.44	29.41	-30.49	-30.47	-30.46	-30.96	-165.71	-166.31	-167.71	-163.43
7602.10	2317.12	1.38	29.59	-29.88	-29.91	-30.10	-30.88	-171.9	-158.94	-163.68	-162.57
7631.60	2326.11	1.60	28.28	-28.81	-29.35	-29.87	-30.60	-140.22	-140.79	-140.16	-130.68
7642.00	2329.28	1.55	29.62	-30.22	-30.87	-31.32	-32.27				-143.54
7787.70	2373.69	1.62	28.88	-29.25	-29.01	-29.50	-30.36	-148.74	-152.08	-159.05	-148.25
7848.40	2392.19	1.62	28.88	-29.47	-29.87	-30.42	-31.48	-122.14	-126.66	-130.17	-130.11
7989.20	2435.11						-164.0	-155.26	-151.75		-133.74

Table S8. Diversity summary results for Mar2x, Riecko Mache and Gonzales sites
 N=number of pollen and spores counted per sample
 S=number of species per sample
 SD=Standing Diversity after range through method and excluding singletons

SD=Standing Diversity, after range-through method

numbers in red indicate sa-

SI=Simpson Index (it was not calculated for samples with counts <80)

MAR 2x Riecito Mache Gonzales

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Table S9. Relative abundances of dry vs wet indicator for Mar2x core. Precipitation preferences of families identified in this study are derived from Punyasena study (S79) of Gentry's 144-transect neotropical plant database.

Family	Precipitation Preferences		Relative Abundance, MAR2x core			
	wet	dry	depth (m.)	depth (ft.)	wet (%)	dry (%)
Arecaceae	1		2080.9	6827.2	59.4	2.6
Olacaceae	1		2081	6827.3	58.4	1.3
Araceae	1		2082.7	6833	37.3	0
Annonaceae	1		2082.7	6833.1	46.1	2.2
Ericaceae	1		2085.2	6841.3	48.5	2.7
Moraceae	1		2088.6	6852.3	38.1	3.1
Convolvulaceae	1		2090.3	6857.8	59.6	1.5
Sapotaceae	1		2090.9	6859.8	68.5	0.8
Passifloraceae	1		2099.2	6887.3	70.7	1.7
Malvaceae	1		2100.7	6892.2	52.5	0
Podocarpaceae	1		2103.1	6900	38.8	1
Fabaceae	1		2103.3	6900.5	60.5	5.9
Euphorbiaceae	1		2105	6906.3	57.5	3.3
Proteaceae	1		2106.8	6912.2	47.9	1.2
Poaceae	1		2108.5	6917.8	54.4	1.5
Podocarpaceae	1		2114	6935.7	59.5	0
Ulmaceae	1		2116.6	6944.3	50.4	0
Myrtaceae	1		2117.8	6948	48.5	0.5
Onagraceae	1		2122.4	6963.1	72.2	1.3
			2123.6	6967.1	82.5	0
			2126	6975	82.7	0
			2127.3	6979.2	78.4	2.6
			2128.8	6984.1	50.2	4.2
			2129.4	6986.1	56.3	1.9
			2129.4	6986.2	74.2	3.1
			2130	6988.2	66.4	4.7
			2131.4	6992.8	59.7	0.8
			2137.4	7012.6	53.6	0.7
			2139.3	7018.8	32	5
			2141.1	7024.5	60.4	2.2
			2143	7030.8	59	0
			2145.4	7038.8	55.3	1
			2149.2	7051.2	45.6	2.5
			2151.9	7060.2	47.7	5.6
			2154.9	7069.9	51.3	3.9
			2155.5	7071.9	70.3	2.2
			2156.1	7073.8	47.9	3.3
			2156.7	7075.8	76.1	1.1
			2159.1	7083.8	74.5	1.4
			2160.1	7087.1	65.3	4
			2163.1	7096.8	68.6	2.9
			2163.5	7098	60	0
			2167.9	7112.4	49	5
			2168.6	7114.7	46.1	4.5
			2169.2	7116.8	61.2	3.1
			2170.2	7120.1	73.8	1.2
			2171.7	7125.1	67.1	2.9
			2173.3	7130.1	65.9	2.3
			2173.6	7131.1	100	0
			2176.1	7139.3	78.6	1
			2177.8	7145	67.6	7
			2178.9	7148.6	76.2	1.4
			2179.9	7151.9	69.1	3.6
			2180.5	7154	72.9	1
			2183.7	7164.5	73.1	3.3
			2193.1	7195.3	45.7	3.5
			2196.5	7206.4	64.6	2
			2225.1	7300.3	44.7	2.2
			2229.2	7313.7	72.9	1.9
			2235.4	7334.1	67.7	7.5
			2241.9	7355.4	67.6	0.7
			2248.4	7376.5	71.6	4.2
			2255.1	7398.5	63.6	1.8
			2266.8	7437.1	46.2	2.3
			2266.9	7437.2	50.7	2.6
			2274.1	7461.1	63	2.4
			2287.6	7505.3	58	3.5
			2290.9	7516.2	67.2	1.5
			2291	7516.3	65	0.5
			2292.2	7520.3	54	2.3
			2294.2	7527	54.2	0.7
			2305.1	7562.7	83	0.7
			2306.7	7568	73.2	0.8
			2308.9	7575.2	56.9	5.5
			2310.6	7580.6	51.8	4.4
			2311.1	7582.5	73.4	0
			2311.7	7584.4	73.9	1
			2313	7588.5	62.7	0.4
			2315.5	7596.9	75.2	1.6
			2316.2	7599.08	49.7	0
			2316.9	7601.3	57.8	0
			2324.7	7627.1	52.2	0.6
			2325.7	7630.2	50.5	3.9
			2326	7631.08	62.2	3.5
			2326.5	7632.9	63.9	0.7
			2329	7641.1	9.2	0
			2338.9	7673.7	96.3	0
			2339.1	7674.1	48.7	0.3
			2342.2	7684.3	98.2	0
			2342.8	7686.3	96.5	0
			2343.2	7687.7	97	0
			2343.3	7688.1	95.9	0
			2343.4	7688.2	91.7	0
			2344	7690.2	96.1	0
			2373.7	7787.6	81.8	0
			2373.8	7788.2	29.1	3.4
			2380.2	7809	47.3	1.1
			2392.2	7848.3	84.6	0
			2392.6	7849.7	28.1	0
			2407.8	7899.5	42.6	1.5
			2421.5	7944.6	52.6	0
			2434.9	7988.5	41.6	0.6
			2436.2	7992.7	74.3	0
			2437.1	7995.8	62.2	0.5
			2437.4	7996.7	40	2.1
			2438.9	8001.6	93.9	0
			2455.8	8057.16	47.5	1.7
			2468.1	8097.5	29.4	0
			2484.1	8150	77.6	0.4
			2492.7	8178	86.2	0

2183.7	7164.5	73.1	3.3
2193.1	7195.3	45.7	3.5
2196.5	7206.4	64.6	2
2225.1	7300.3	44.7	2.2
2229.2	7313.7	72.9	1.9
2235.4	7334.1	67.7	7.5
2241.9	7355.4	67.6	0.7
2248.4	7376.5	71.6	4.2
2255.1	7398.5	63.6	1.8
2266.8	7437.1	46.2	2.3
2266.9	7437.2	50.7	2.6
2274.1	7461.1	63	2.4
2287.6	7505.3	58	3.5
2290.9	7516.2	67.2	1.5
2291	7516.3	65	0.5
2292.2	7520.3	54	2.3
2294.2	7527	54.2	0.7
2305.1	7562.7	83	0.7
2306.7	7568	73.2	0.8
2308.9	7575.2	56.9	5.5
2310.6	7580.6	51.8	4.4
2311.1	7582.5	73.4	0
2311.7	7584.4	73.9	1
2313	7588.5	62.7	0.4
2315.5	7596.9	75.2	1.6
2316.2	7599.08	49.7	0
2316.9	7601.3	57.8	0
2324.7	7627.1	52.2	0.6
2325.7	7630.2	50.5	3.9
2326	7631.08	62.2	3.5
2326.5	7632.9	63.9	0.7
2329	7641.1	9.2	0
2338.9	7673.7	96.3	0
2339.1	7674.1	48.7	0.3
2342.2	7684.3	98.2	0
2342.8	7686.3	96.5	0
2343.2	7687.7	97	0
2343.3	7688.1	95.9	0
2343.4	7688.2	91.7	0
2344	7690.2	96.1	0
2373.7	7787.6	81.8	0
2373.8	7788.2	29.1	3.4
2380.2	7809	47.3	1.1
2392.2	7848.3	84.6	0
2392.6	7849.7	28.1	0
2407.8	7899.5	42.6	1.5
2421.5	7944.6	52.6	0
2434.9	7988.5	41.6	0.6
2436.2	7992.7	74.3	0
2437.1	7995.8	62.2	0.5
2437.4	7996.7	40	2.1
2438.9	8001.6	93.9	0
2455.8	8057.16	47.5	1.7
2468.1	8097.5	29.4	0
2484.1	8150	77.6	0.4
2492.7	8178	86.2	0

Table S10. P2 core TEX86 values, BIT indices and SST estimates following the calibration of Kim et al (S85) and Liu et al (S86).

One standard deviation for the temperature estimation is 3.6 C (using Liu calibration (S86))

Elevation (m)	Age	TEX86	BIT	SST (Kim et al.)	SST (Liu et al.)
123	Late Paleocene	0.71	0	28.6	27.6
129.9	Late Paleocene	0.7	0	28.0	27.2
157	Late Paleocene	0.68	0.05	27.1	26.4
181.25	Late Paleocene	0.65	0	26.0	25.5
189.36	Late Paleocene	0.69	0	27.6	26.8
229.86	Late Paleocene	0.68	0	27.1	26.5
238	Late Paleocene	0.78	0	31.3	29.6
277.5	Late Paleocene	0.71	0	28.3	27.3
292.78	Late Paleocene	0.71	0.05	28.5	27.5
301.95	Late Paleocene	0.71	0.05	28.6	27.6
309.62	Late Paleocene	0.73	0	29.4	28.2
316.62	Early Eocene	0.75	0.05	30.0	28.6
336.4	Early Eocene	0.75	0.1	29.9	28.6
359.24	Early Eocene	0.79	0.12	31.8	29.9
372.74	Early Eocene	0.7	0.06	27.9	27.1
390.29	Early Eocene	0.83	0.09	33.0	30.8
404.3	Early Eocene	0.87	0.11	34.4	31.6
424.1	Early Eocene	0.84	0.13	33.5	31.1