doi: 10.5027/andgeoV41n3-a02

# Chronology and geology of an Early Miocene mammalian assemblage in North of South America, from Cerro La Cruz (Castillo Formation), Lara state, Venezuela: implications in the 'changing course of Orinoco River' hypothesis

\*Ascanio D. Rincón¹, Andrés Solórzano¹, Mouloud Benammi², Patrick Vignaud², H. Gregory McDonald³

- <sup>1</sup> Laboratorio de Paleontología, Centro de Ecología, Instituto Venezolano de Investigaciones Científicas (IVIC), Caracas, Venezuela. paleosur1974@gmail.com; solorzanoandres@gmail.com
- <sup>2</sup> Institut de Paléoprimatologie, Paléontologie Humaine, Evolution et Paleonviroments, UMR CNRS 7262-INEE, Université de Poitiers, France.
  - mouloud.benammi@univ-poitiers.fr; patrick.vignaud@univ-poitiers.fr
- <sup>3</sup> Museum Management Program, National Park Service, 1201 Oakridge Drive, Fort Collins, Colorado 80525, U.S.A. greg mcdonald@nps.gov
- \* Corresponding author: paleosur1974@gmail.com

ABSTRACT. In general the geology of paleontological sites in Venezuela is poorly known. With the purpose of improving this knowledge we describe the geology of the Castillo Formation (Late Oligocene to Early Miocene) at Cerro La Cruz locality, in Lara state, Venezuela, that contains several records of vertebrate and invertebrate fauna. Lithologically, the Cerro La Cruz sequence is composed by alternating packages of siliciclastic and carbonate sediments, with a predominance of mudstone. The paleoenvironment is inferred as a mainly near-shore marine complex that could be associated with regressive and transgressive phases. Nevertheless, into the middle part of the Cerro La Cruz outcrops two levels containing at least six mammalian remains were found, confirming the early continental mammal assemblage in Venezuela. The continental vertebrate assemblage includes Xenarthra, Notoungulata and Litopterna mammals, fresh water fishes, the turtle Chelus, wood and leaves, allows us to interpret the paleoenvironment of the continental episodes of this locality as a mosaic composed of an humid forest and lowland savanna. Four isotopic dates using Strontium (87Sr/86Sr), from the Cerro La Cruz locality were obtained, which bracket of the sediments to between 17.21 to 19.27 Ma, confirming an Early Miocene age for this locality. Biochronologically, the mammalian assemblage so far recovered does not permit to assign this fauna to any of the recognized South American Land Mammal Ages, however the isotopic ages suggest affinities with a Pansantacrucian mammalian sub-cycle. Our interpretation of the geology at Cerro La Cruz shows no evidence for the presence of a major river system crossing over that zone during the Early Miocene and does not support the hypothesis of the possible draining of a 'Proto-Orinoco' river into Maracaibo or Falcón basin during the Early Miocene.

Keywords: Early Miocene, Pansantacrucian, Castillo Formation, 'Proto-Orinoco' evolution, Venezuela.

RESUMEN. Cronología y geología de una asociación de mamíferos del Mioceno Temprano en el Norte de América del Sur, cerro La Cruz (Formación Castillo), estado Lara, Venezuela: implicaciones para las hipótesis del 'cambio del curso del río Orinoco'. En general los estudios geológicos detallados de las localidades paleontológicas de Venezuela son escasos. Con la finalidad de mejorar este conocimiento, se describe la geología de la Formación Castillo (Oligoceno Tardío a Mioceno Temprano) en afloramientos localizados en cerro La Cruz, estado Lara, Venezuela, la cual contiene reportes de una diversa fauna de vertebrados e invertebrados. Litológicamente la secuencia observada en cerro La Cruz está compuesta por una intercalación de paquetes siliciclásticos y carbonáticos, con predominio de lutita. El paleoambiente inferido para esta localidad es principalmente de un complejo marino-costero asociado a fases transgresivas y regresivas. Sin embargo, hacia la parte media de la sección del cerro La Cruz se reportan dos estratos portadores de al menos seis restos de mamíferos continentales, los cuales conforman la más diversa y antigua asociación faunística de Venezuela. Las asociaciones de vertebrados continentales que incluyen mamíferos de las familias Xenarthra, Notoungulata y Litopterna, peces de agua dulce, una tortuga del género Chelus, además de restos de madera y hojas, nos permiten interpretar la existencia de un mosaico de ambientes compuesto por bosques húmedos y sabanas bajas. También se presentan cuatro fechamientos isotópicos realizados sobre la base del método del Estroncio (87Sr/86Sr), en la localidad de cerro La Cruz, arrojando edades comprendidas entre 17,21 y 19,27 Ma, lo que confirma una edad Mioceno Temprano para esta localidad. Biocronológicamente la asociación de mamíferos no permite referirlo a ninguna Edad Mamífero de America del Sur, sin embargo, las edades isotópicas sugieren afinidad al subciclo mamaliano del Pansantacrucense. Nuestra interpretación de la geología de cerro La Cruz no muestra evidencias de la presencia de un gran río cruzando esta zona durante el Mioceno Temprano, y no es consecuente con la hipótesis del posible drenaje de un ancestro del río Orinoco hacia las cuencas sedimentarias de Maracaibo o Falcón durante el Mioceno Temprano.

Palabras clave: Mioceno temprano, Pansantacrucense, Formación Castillo, Evolución del 'Proto-Orinoco', Venezuela.

#### 1. Introduction

The Early Miocene continental mammal fauna is poorly known in Venezuela, and has not been recorded from Colombia, Brazil, or Trinidad. The first record of a continental mammal from this age in Venezuela was Xenstrapotherim christi from the Chaguaramas Formation (Early Miocene sensu Wesselingh and Macsotay, 2006) (Stehlin, 1928). Later Collins (1934) reported Pseudoprepotherium venezuelanum from the Río Yuca Formation (Early Miocene sensu Macsotay et al., 1995). The third terrestrial mammal, from the Quiamare Formation (Early to Late Miocene), Boreostemma venezolensis was reported by Simpson (1947). More recently (Sánchez-Villagra et al., 2004; Weston et al., 2004; O'Leary, 2004) reported another Early Miocene locality in the Castillo Formation with fossil mammals.

None of these localities has been dated using radiometric methods so their exact age is unclear. For example the age of the Río Yuca Formation has been determined based on palynological data (Macsotay *et al.*, 1995), and the proposed ages of the Chaguaramas (Fasola *et al.*, 1985), Quiamare and Castillo Formation (Wheeler, 1960, 1963; Lorente, 1987) are based on foraminifera and invertebrate data.

The first study of the faunal assemblage in the Castillo Formation was the molluscan fauna (Wheeler, 1960) from the Cerro Castillo locality. Additional

studies by several authors (Sánchez-Villagra *et al.*, 2000; Dahdul, 2004; Aguilera and Rodríguez de Aguilera, 2004a y b; Sánchez-Villagra *et al.*, 2004; Brochu and Rincón, 2004; Weston *et al.*, 2004; O'Leary, 2004) have reported more or less thirty new faunal records for this formation, seven of which are mammals, but only two of these are terrestrial mammals. Most of these taxa are from a locality named Cerro La Cruz, Lara state. Previously at this locality Sánchez-Villagra *et al.* (2000, 2001) interpreted the depositional environment as near-shore to shallow marine for the base to 87 m of the exposed section.

Our research has documented 367 m of stratigraphic section that includes different sedimentary units from those previously reported. In this paper we describe these new sedimentary units of the Castillo Formation and present new strontium isotope (87Sr/86Sr) dates. In addition, we discuss the age and paleoenvironment of the exposures of the Castillo Formation at Cerro La Cruz and it implications in a South American context, specially taking into account the hypothesis of the 'changing of course of the Proto-Orinoco River' in northern of South America.

# 2. Reviews of paleontological, geological settings and age of Castillo Formation

The Castillo Formation was deposited in the Tertiary Falcón Basin, which extends into the states

of Falcón and parts of Zulia, Lara and Yaracuy in northwestern Venezuela. Wheeler's (1960) original description of the Castillo Formation was based on outcrops at Cerro Castillo, located about 27 km to the southwest of Dabajuro, Buchivacoa Municipality, Falcón State.

The Falcón Basin is situated within the Caribbean-South American Plates boundary area. The framework of the basin is the result of the polyphase tectonics and is rather complex (Baquero et al., 2009). It involves Late Cretaceous to Middle Eocene emplacement of the Lara Nappes followed by Late Eocene to Early Miocene tectonic collapse and graben formation, and by Middle Miocene inversion and out of sequence thrusting (Baquero et al., 2009). The sedimentary record within the Falcón Basin has been almost continuous since Late Eocene time, except for three angular and/or erosional unconformities of some regional extent (Audemard, 1993, 2001). During the Oligocene and Miocene, the marine Falcón Basin was an elongate east-west trending depression bordered by topographic highs to the south, west and north, and formed the western end of the larger Bonaire Basin, at present the offshore portion of the Venezuelan Coast Range (Muessig, 1984; Audemard, 1993). Throughout the Middle-Late Miocene, NW-SE compression resulted in folding, tectonic inversion, and exposure of a nearly continuous record of sedimentation from the Eocene to the Pliocene (Audemard, 1993, 2001). The Castillo Formation during the Oligo-Miocene conforms to the northwest to southeast edge of the Falcón Basin, and its outcrops are found in a wide semicircular area that extends through the states of Falcón and Lara (Wheeler, 1960).

Wheeler (1960) suggested that the Formation was deposited in environments ranging from shallow water to brackish facies, with local continental facies. Based on the foraminifera and invertebrates, Wheeler (1960) estimated that the age of the Castillo Formation ranged from Late Oligocene in the north to Early Miocene in the south of the Falcón Basin. Lorente (1987) identified from drill hole TIG-141X (Castillo Formation North in Lara state, near type section) a sporomorph assemblage which was assigned to the Magnastriatites-Cicatricosisporites dorogensis zone (Late Oligocene) in the lower part of the formation, and the *Verrutricolporites* zone, which is indicative of an Early Miocene age in the upper part of the formation. Later, Sánchez-Villagra et al. (2000), based on the foraminifera assemblage, considered the entire Castillo Formation to be Early Miocene in age following the zonation of Cati et al. (1968). Johnson et al. (2009) studied the scleractinian reef coral assemblages from four different units in the northwestern portion of the Falcón Basin of Venezuela, including one from the Castillo Formation at Cerro Guariro (northern part of Formation) locality. The coral remains found in this assemblage included distinctive Oligocene genera such as Antiguastrea and Diploastrea, and suggest that at least at Cerro Guariro the Castillo Formation is Late Oligocene in age. López and Brineman (1943) proposed that the Castillo Formation in the south of Baragua Sierra is a typically sublittoral shallow water marine environment, with an Early Miocene age for this southern part of Castillo Formation.

From 87 m of clayey marls interbedded with numerous thin hardground units Sánchez-Villagra et al. (2000) reported the first vertebrate faunal assemblage from outcrops of the Castillo Formation at the Cerro La Cruz locality, in Lara State. These authors report a fauna composed of one turtle, one crocodile, two whales (Odontoceti), three sharks, along with a palm fruit, 20 molluscan species, and one crab. Subsequent studies carried out at the Cerro La Cruz locality (Sánchez-Villagra et al., 2001; Dahdul, 2004; Aguilera and Rodríguez de Aguilera, 2004a and b; Sánchez-Villagra et al., 2004; Weston et al., 2004; O'Leary, 2004; Brochu and Rincón, 2004; Aguilera et al., 2010; Aguilera and Lundberg, 2010; Sánchez-Villagra et al., 2010; Rincón et al., 2010a and b; Aguilera et al., 2013) resulted in the recognition of a diverse vertebrate assemblage.

Additional records of vertebrate fauna from the Castillo Formation consist of a new genus and species of gharials, Siquisiquesuchus venezuelensis, which holotype was found in the vicinity of Siquisique Town, in the eastern of the Sierra de Baragua, Lara State (Brochu and Rincón, 2004), and fragmentary turtle and crocodile remains from the northwest of Bariro Town, northern part of the Castillo Formation, in Falcón State (Sánchez-Villagra et al., 2010). The enigmatic Pyrotheria Proticia venezuelensis, was described by Patterson (1977) from the Trujillo Formation (Paleocene-Eocene). The Trujillo Formation represents a turbidite system, with a micromolluses fauna that suggests a palaeobathymetry of 2-5 km, in a deep epicontinental slope environment (Mactosay et al., 1989). Several authors (Sánchez-Villagra et al., 2000, 2001, 2010; Bond and Gelfo, 2010)

considered that the provence of *P. venezuelensis* should be put into a question, suggesting that this fossil was collected into Miocene rocks of the Castillo Formation instead in the Paleocene-Eocene rocks of the Trujillo Formation.

An updated vertebrate faunal list for the Cerro La Cruz locality is provided in Table 1, including the recent records of terrestrial mammals (Rincón *et al.*, 2010a and b).

## 3. The Geology of Cerro La Cruz

Cerro La Cruz is located around 78 km to the southeast of the type section of the Castillo Formation, close to La Mesa village, on the south flank of Sierra de la Baragua (10°22'05.16"N and 70°04'14.45"W), in the Pedro Leon Torres Municipality, Lara State, 20 km north of Carora town (Fig. 1). It has a well-exposed stratigraphic sequence of at least 367 m (Fig. 2) that was described here. The sequence consists predominately of mudstone, interbedded with limestone and sandstone, with local units of limonite and conglomerate. From the base to top the following units are recognized:

• Unit A. Total thickness is 96 m. It consists of alternating layers of mudstone, sandstone and conglomerate with the mudstone as the most abundant lithology (Fig. 3). They are bioturbated, motley grey with reddish and yellowish bands (jarosite). The individual strata thicknesses vary between 1-20 m. The sandstone presents light colors; with thicknesses varying between 30-200 cm. Mineralogically the sandstone is essentially composed of quartz, with a large grain size (0.8-0.1 mm), conglomeratic lenses, bad to moderately well-sorted to the base, and with fine grains, and well packed to the top. The thick sand layers were observed to be formed from moderately to well-rounded quartz fragments and metamorphic rocks, and at the base have local reworked oyster fragments. Ripple marks, low angle cross-bedded and parallel stratification are also observed in sandstone. The unit exhibits, in general, the presence of oxidation levels that increase towards the top of the strata. The bioturbation has a grade of intensity one (Droser and Bottjer, 1986) and two burrowed ichnofabrics are identified in this unit, Thalassinoides (Fig. 4a) and Ophiomorpha. The unit is poorly fossiliferous, but in the middle part contains some wood fragments and toward the top organic matter. Toward the base Ostrea

remains are also present. The inferred depositional environment for this unit is near-shore marine with limited clastic terrigenous influx toward the top.

• Unit B. Total thickness 118 m. It consists of alternating mudstone, bioclastic limestone and sandstone (Fig. 3). The color of the mudstone ranges from light grey to lead-grey, is generally fossiliferous and bioturbated, with some iron oxide nodules toward the base. There is locally abundant organic matter. The thickness of the layers varies from 2.5-11 m. The limestone is bioclastic, consisting of fragments of invertebrates that are sometimes reworked, and ranges in color from orange to purple. There has been some compaction, the thicknesses ranges between 15-50 cm. Dunham (1962) classified them as: packstone, mudstone and wackstone with packstone predominant (with micritic matrix). Some limestone grades laterally into sandstone. The few sandstone present are restricted to the base of unit, have thick layers and large grains (0.8-0.1 mm), with decreasing size grain toward the top, with conglomerate levels, composed of quartz fragments and metamorphic rock remains toward the base. Bioturbation is conspicuous towards the top, and has a grade of intensity one to two (Droser and Bottjer, 1986), of the Ophiomorpha and Thalassinoides ichnofacies. Wave ripple marks are conspicuous toward the base (Figs. 4b and c).

This unit is highly fossiliferous and preserves an abundance of invertebrate fossils including: Anadara sp., Chionopsis tegulum, Clementia dariana dariana, Cyclinella cyclica, Liromissus quirosensis, Trachycardium sp., Glyptoactis quirosana, Ostrea sp., Pecten sp., Saccella gracillima, Architectonica nobilis, Cirsotrema sp., Conus sp., Melongena cf. M. venezuelana, Turritella larensis, Turritella venezuelana, Turritela sp., Dentalium bocasense, abundant crab remains Calappa larensis, Eriosalchila rathbunae, Palaeopinnixa perornata, Portunus aff. P. oblongus and some Cnidaria indet. Toward the top many fish species (otolith based) are conspicuous along with shark and ray teeth. Toward the base indeterminate turtle remains and wood is present.

Following the Standard Microfacies Model (SMF) of Wilson (1975) and Flügel (2004), the presence of bioclastic wackestone/packstone found in this unit (some of them composed of well-preserved skeletal shells and fragments) is interpreted as having been deposited in a shelf lagoon with

TABLE 1. UPDATED VERTEBRATE FAUNAL LIST AT CERRO LA CRUZ, CASTILLO FORMATION.

Class	Order	Family	Genera	Taxa
Chondrichthyes	Lamniformes	Otodontidae	Megaselachus	M. megalodon <sup>1</sup>
	Orectolobiformes	Ginglymostomatidae Nebrius		Nebrius delfortriei <sup>2</sup>
	Carcharhiniformes	Hemigaleidae	Hemipristis	Hemipristis serra <sup>1</sup>
		Carcharhinidae	Carcharhinus	cf. C. obscurus <sup>1</sup>
				cf. C. perezi¹
				C. falciformis <sup>1</sup>
				C. plumbeus <sup>1</sup>
		Myliobatidae	Myliobatis	Myliobatis sp. <sup>1</sup>
		•	Rhinoptera	Rhinoptera sp. <sup>1</sup>
Actinopterygii	Characiformes	Serrasalmidae	Colossoma	C. macropomum <sup>1</sup>
			Mylossoma	Mylossoma sp.¹
	Siluriformes	Ariidae	Bagre	B. protocaribeanus <sup>3</sup>
			Cantarius	C. nolfi <sup>3</sup>
	Perciformes	Sciaenidae	Ctenosciaena	aff. C. gracilicirrhus <sup>1</sup>
			Equetus	Equetus davidandrewi <sup>1</sup>
			Paralonchurus	P. schwarzhansi <sup>1</sup>
				P. trinidadensis <sup>1</sup>
			Plagioscion	Plagioscion marinus <sup>1</sup>
			Protosciaena	Protosciaena neritica <sup>1</sup>
			Larimus	Larimus henrici¹
		Sphyraenidae	Sphyraena	Sphyraena sp.¹
		Scombridae	Achanthocybium	Achanthocybium sp. <sup>1</sup>
Sauropsida	Crocodilia	Gavialidae	Siquisiquesuchus	S. venezuelensis. <sup>4</sup>
			7	Tomistominae indet. <sup>5</sup>
		Alligatoriidae		Alligatoriidae, indet. <sup>2</sup>
	Testudines	Trionychidae		Trionychidae, indet. <sup>6</sup>
		Podocnemidae	Bairdemys	Bairdemys sp.8
		Chelidae	Chelus	Chelus sp. <sup>2</sup>
Mammalia	Cetartiodactyla	Squalodontidae	Prosqualodon	aff Paustralis <sup>7</sup>
		Squalodelphinidae	Notocetus	aff. N. vanbenedini <sup>9</sup>
		Iniidae	110100011115	Iniidae, indet. <sup>6</sup>
		Platanistidae		Platanistoidea, indet. <sup>6</sup>
	Sirenia	1 Manistrate		Sirenia, indet. <sup>6</sup>
	Xenarthra	Orophodontidae		Orophodontidae, Indet. <sup>10</sup>
		Pampatheriidae	Scirrotherium	Scirrotherium sp. 10
		- umpumernuuc	zon omer min	Phyllophaga indet. <sup>6</sup>
	Notoungulata	Astrapotheriidae		Astrapotheriinae, IndetA <sup>6</sup>
	rotounguiata	Astrapoulcinuae		Astrapotheriinae, IndetB <sup>6</sup>
	Litontarna			•
	Litopterna			Litopterna, indet. <sup>10</sup>

<sup>&</sup>lt;sup>1</sup> Aguilera and Lundberg, 2010; <sup>2</sup> Rincón *et al.*, 2010b; <sup>3</sup> Aguilera *et al.*, 2013; <sup>4</sup> Brochu and Rincón, 2004; <sup>5</sup> Sánchez-Villagra *et al.*, 2001; <sup>6</sup> Sánchez-Villagra *et al.*, 2000; <sup>8</sup> Sánchez-Villagra *et al.*, 2004; <sup>9</sup> O'Leary, 2004; <sup>10</sup> Rincón *et al.*, 2010a.

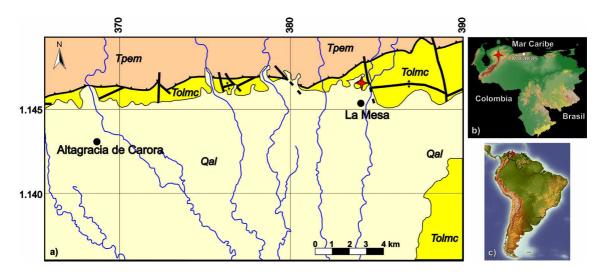


FIG. 1. Location and geological setting of Cerro La Cruz, Castillo Formation, Lara state, Venezuela. Red star shows the location of the analyzed section. a. Geological map of Cerro La Cruz Locality, modified from Martínez and Valletta (2008); b. Regional;
 c. Continental map of the location of the described section. Abbreviations: Tpem= Matatere Formation (Eocene), Tolmc= Castillo Formation (Early Miocene), Qal= Alluvial (Quaternary).



FIG. 2. Cerro La Cruz landscape with the outcrops of Castillo Formation.

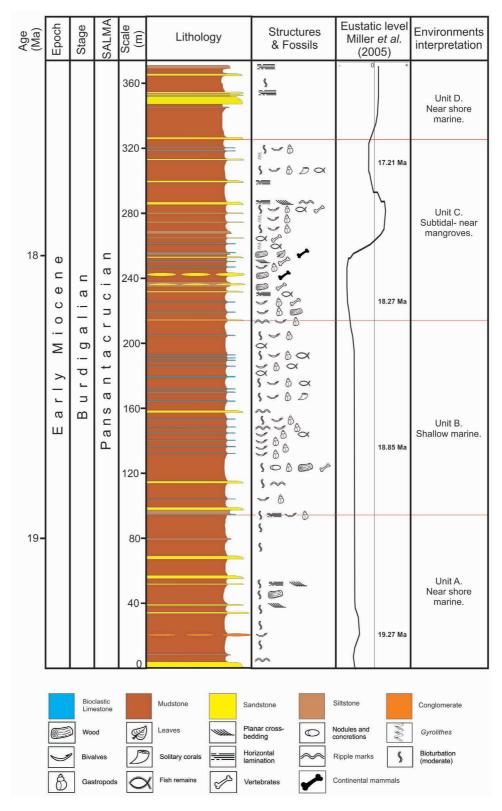


FIG. 3. Stratigraphic column of Cerro La Cruz, Castillo Formation, Lara State.



FIG. 4. Sedimentary structures in Cerro La Cruz, Castillo Formation, Lara State. a. Thalassinoides; b. Oblique view of wave ripple marks; c. Lateral view of wave ripples marks; d. *Gyrolithes*.

low energy water circulation (SMF No. 8). The bioclastic mudstone is interpreted as deposited in shallow water with open circulation close to wave base? (SMF No. 9).

The high diversity of invertebrate remains is conspicuous in this unit. They are better preserved in the mudstone layers whereas in bioclastical limestone they are sometimes reworked. In mudstone the low degree of disarticulation of the bivalves and gastropods, also suggests a low energy conditions, with a short high energy interlude during which the fauna was smothered and buried by sediments, with minimal post-mortem disturbance. Moreover the Thalassinoides-Ophiomorpha ichnofacies assemblage could be interpreted as having developed on subtidal substrates under well aerated agitated shallow waters (Verde, 2002), which is consistent with the presence of wave ripple marks (Figs. 4b and c). The presence of many oxidation levels is interpreted as caused by short periods of subaerial exposure. The poor sandstone development indicates a limited contribution of continental clastic sediments. In general this configuration is consistent with shallow water, with well aerated agitated waters, that is probably slightly deeper towards the top, limiting the development of ripple marks and clastic input, and encouraging the presence of fishes.

• Unit C. Total thickness is 110 m. It consists of alternating mudstone, sandstone and bioclastic limestone, with mudstone being most conspicuous (Fig. 3). This unit includes the top of Cerro La Cruz hill. The mudstone ranges between light grey, lead grey and dark gray, is generally fossiliferous, mostly toward the base, and is bioturbated mostly toward the top. Thin iron oxidation levels (hematite and goethite) are common and interbedded between mudstone layers. Subarkose sandstone is the second predominant lithology (Dott, 1964) and is composed of medium to fine grains (0.6-0.1 mm)

with muscovite, altered plagioclase, and quartz. The grains are finer toward the top, generally with subrounded and moderately well-sorted grains. They are moderately to well packed, and range from 30-200 cm in thickness. The contact between grains is punctual and always surrounded by the matrix. Cement is present and in part composed of iron oxide (limonite). Toward the base the sandstone can have lenticular bedding, but in general the strata are tubular. Ripple marks, parallel lamination and stratification, and planar cross-bedding are common. The limestone is bioclastic, consisting of many fragments of invertebrates sometimes reworked specially toward the base, which range in color from orange to purple, being more abundant toward the base. There has been some compaction, the thicknesses range between 20-35 cm. Dunham (1962) classified them as: grainstone (with spar cement), packstone, and wackstone, with packstone being predominant (with micritic matrix). This unit presents a highly lateral variation, in some cases the bioclastic limestone will laterally transition into sandstone and vice versa.

In general, most of the published fauna from Cerro La Cruz (Sánchez-Villagra et al., 2004; Aguilera et al., 2010; Sánchez-Villagra et al., 2010) comes from this unit. The unit preserves an abundance of marine invertebrate fossils including: Anadara mirandana, Anadara cf. A. inutilis, Chionopsis tegulum, Clementia dariana dariana, Trachycardium sp., Glyptoactis quirosana, Ostrea sp., Pecten sp., Saccella gracillima, Architectonica nobilis, Conus sp., Melongena cf. M. venezuelana, Turritella larensis, Turritella venezuelana, Turritella montanitensis, Turritela sp., Dentalium bocasense, and abundant indeterminate crab remains, mostly present in the bioclastic limestone and mudstone. Are also present are many marine fish species (mostly based on otoliths), ray and shark teeth, while toward the top of the unit in the sandstone bed Cetartiodactyla remains, Squalodontidae, aff. Prosqualodon australis (Sánchez-Villagra et al., 2000), and Squalodelphinidae aff. Notocetus vanbenedini (O'Leary, 2004) have been reported. In addition to the marine fauna found in this unit, in two level located toward the base of this unit Rincón et al. (2010a y b) reported a varied 'continental or fresh water' fauna (housed at the paleontological collection in the Instituto Venezolano de Investigaciones Científicas (IVIC), Caracas, Venezuela)

represented by an astragalus (IVIC-P-1829) with the typical spool-like astragalar body characteristic of a Litoptern (Fig. 5a); a mandible (IVIC-P-1830) with the external sulcus on m1 lacking, but present in m2, characteristic of members of the Astrapotheriinae (Fig. 5b); a partial imbricating osteoderm (IVIC-P-1827) that is relatively thin with a small number of well-spaced follicular pits connected by a distinct channel, typical of the pampathere, Scirrotherium (Fig. 5c); and a partial mandible (IVIC-P-1828) with the alveolus of the first molariform, and the second to fourth molariforms. composed of a very thin outer layer of cement (0.5 mm), a thicker internal layer of osteodentine (6.3 mm) and a core of vasodentine (1.3 mm), diagnostic of the sloth family Orophodontidae sensu Hoffstetter, 1969 (Fig. 5d). Additional 'continental or fresh water' fauna mentioned by Rincón et al. (2010a and b) consist of Alligatoriidae indet., turtle remains of Chelus sp. (IVIC-P-2136), and some Testudine indet. In the lenticular sandstone and sandy siltstone associated with continental vertebrate fauna several teeth of fresh water Serrasalminae (Colossoma sp. and Mylossoma sp.), previously reported from this locality by Dahdul (2004) were recovered along with common plant leaves that were found in association with the continental mammal remains.

Some levels are bioturbated, especially toward the middle and top, and have a grade of intensity one to two (Droser and Bottjer, 1986). Ophiomorpha, Thalassinoides and Gyrolithes ichnofacies are recognized, with the last two being conspicuous. The Gyrolithes ichnofacies is restricted to a middle upper part of the unit, (Fig. 4d) and is commonly reported from brackish-water mudstone, siltstone and sandstone, in marginal-marine settings, and are also associated to intertidal zones, near mangrove areas (mud flats and channels, Netto et al., 2007). The presence of Gyrolithes combined with abundant wood remains (some associated with oysters in living position) were observed in the middle and upper part of the unit and reflects episodes of brackish-water probably near mangrove areas or estuarine environments. The near-shore marine complex environments are dominant in this unit, but the mammal assemblage is found only in two levels from the middle part of this unit indicating the presence of this continental environment only for a short duration.

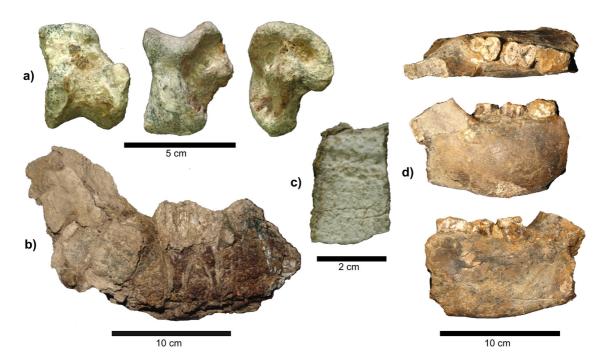


FIG. 5. Mammal fossils from Cerro La Cruz, Castillo Formation. a. Astragalus of Litoptern (IVIC-P-1829); b. Mandible of Astrapotheriinae (IVIC-P-1830); c. Partial imbricating osteoderm of *Scirrotherium* (IVIC-P-1827); d. Partial mandible of Orophodontidae (IVIC-P-1828).

• Unit D. Total thickness is 43 m. It mainly consists of alternating mudstone and sandstone (Fig. 3). The color of the mudstone ranges from light grey to mottled-grey, and is not fossiliferous, generally it contains organic matter, and some iron oxide nodules are present toward the base. The thickness of the layers varies from 0.5-18 m. The sandstone presents light colors to mottled; with thicknesses varying between 1-2 m. Mineralogically it is essentially composed of quartz, with scarce muscovite, thin grain size (0.25-0.125 mm), that are moderately to well sorted, rounded, and subspherical. In the sandstone are conspicuous floating rounded pebbles (up to 4 cm), consisting of quartz and metamorphic rock fragments reflecting rapid changes in energy level. In general the sandstones present parallel lamination. The inferred deposition environment for this unit is near-shore marine.

#### 3.1. General settings

Gypsum is abundant in most of the sequence, but this cannot be considered as evidence for an evaporitic environment because its location is restricted to oblique and vertical joints and fractures and is typical of secondary mineralization.

Towards the top of the section, the strata analyzed were in tectonic and in discordant contact with the underlying rocks of the Matatere Formation (not Misoa Formation sensu Sánchez-Villagra et al., 2000; Fig. 1a) of Eocene age, that consist of a type flysch sequence, a monotonous alternation of sandstone and shale with conglomeratic levels and blocks (olistolit and olistostrome) of different lithologies embedded in the sequence. The contact between Matatere and Castillo Formations in the vicinity of Cerro La Cruz is a thrust fault, caused by compressive stress associated with tectonic inversion of the Falcón Basin which started in the Middle Miocene (Baquero et al., 2009; Urbani and Mendi, 2011). This thrust fault relationship documents changes in subsidence and uplift during the evolution of the Falcón Basin (Baquero et al., 2009). Lower contacts are not observed in the outcrops. The section at Cerro La Cruz is located on the northeast flank of an elongated dome on a hinge line oriented N65E. In general, the strata analyzed presents an orientation of N65E with a dip slope varying between 20° and 40°N. In general, outcrops

of the Castillo Formation in Baragua Sierra, are highly deformed, with folding axes subparallel to the thrust front.

## 3.2. 87Sr/86Sr Dating

Precise and detailed data on the variability of strontium isotopes with time in the world's oceans is known for a significant part of Phanerozoic time (Howarth and McArthur, 1997), and local curves for specific stratigraphic sections are being increasingly used to infer absolute ages (Graham *et al.*, 2000). For the middle Cenozoic in particular, the strontium isotope sea-water curve changes rapidly with time, and so is particularly suitable for geochronology (Graham *et al.*, 2000). In this time interval, resolution of stage boundaries is better than 0.5 my (Oslick *et al.*, 1994; Howarth and McArthur, 1997), making the method a powerful tool for improving correlation between biostratigraphic and chronostratigraphic timescales (Graham *et al.*, 2000).

We analyzed the remains of four invertebrates from Cerro La Cruz beds on the South flank of the Sierra de la Baragua, Lara State. We used a Mass Spectrometry Finnigan MAT-262 to determinate the isotopic relationship of the Strontium (87Sr/86Sr), with a Standard laboratory value NBS987=0.710240±28, n=422. This analysis was made by the Laboratorio Universitario de Geoquímica Isotópica (LUGIS) at the Universidad Nacional Autónoma de México, México. Results are shown in the Table 2, and provide an age range between 17.21 and 19.27 Ma.

#### 4. Discussion

#### 4.1. Age

The isotopic age obtaining by the Strontium (87Sr/86Sr) relationship shows that sediments at the

Cerro La Cruz locality were deposited between 17.21 (+0.41/-0.49) to 19.27 (+0.64/-0.53) Ma, and allows to us to infer that the sediments in the stratigraphic section at the Cerro La Cruz to be Early Late Miocene in age (Burdigalian age), including the Magnetic Chrons C5Cr, C5Dn, C5Cn, C5Dr and C5En (Cande and Kent, 1995). This age corroborates the previously estimated age proposed by several authors based on foraminifera and invertebrates (López and Brineman, 1943; Wheeler, 1960; Sánchez-Villagra et al., 2000). In addition, it allows a refinement the age of the vertebrate fauna found in this locality, confirming that the Cerro La Cruz Fauna represents the oldest mammal assemblage of Venezuela, as proposed by several authors (Sánchez-Villagra et al., 2000, 2010; Bond and Gelfo, 2010), and hence the enigmatic *Proticia venezuelensis* would be highly probably from Castillo Formation at Quebrada AguaViva, located about 5 km northwest of Cerro La Cruz.

Within the context of the South American Land Mammal Ages (SALMA), the isotopic values obtained in the stratigraphic section of Cerro La Cruz allows to us to assign a post-Colhuehuapian SALMA (~20 Ma sensu Ré et al., 2010), and pre-Santacrucian SALMA (17.8 to ~16.0 Ma sensu Marshall et al., 1986) age to the fauna. Nonetheless the Cerro La Cruz locality is chronologically closer in time to the Santacrucian and 'Pinturan' SALMA. The type area for the 'Pinturan' SALMA is the northeast of Santa Cruz Province, Argentina and has been dated to be between 16.5 and 17.5 Ma (Kramarz et al., 2010). However, Dunn et al. (2012) based on the section at the Gran Barranca, Central Patagonia, Argentina, proposed that the 'Pinturan' SALMA is bracketed between 19.04 and 18.62 Ma, so the maximum duration for the 'Pinturan' is between 19.04-16.5 Ma (Dunn et al., 2012). Otherwise the Santacrucian SALMA age is usually considered to be between 16.3-17.5 Ma (Flynn and Swisher, 1995) or 16-17.8 Ma (Marshall et al., 1986).

TABLE 2. RELATIONSHIP OF THE STRONTIUM ISOTOPES (87Sr/86Sr) IN THE CERRO LA CRUZ SAMPLES.

Sample	<sup>87</sup> Sr/ <sup>86</sup> Sr	1 sd*	2 SE(M)	n	Туре	Ma	Ma (+/-)	Depth (m)
Cast-58	0.708667	30	4	57	Turritela sp.	17.21	0.41/0.49	305
Cast-38	0.709	25	7	54	Shell	18.27	0.42/0.37	230
Cast-6	0.70853	38	5	58	Ostrea sp.	18.85	0.70/0.61	138
Cast-B25	0.708502	28	4	58	Ostrea sp.	19.27	0.64/0.53	20

n= Value of relationship made by series; 1 sd=1 Standard deviation, 2 SE(M) = 2sd/root n.

In contrast Croft *et al.* (2004, 2007) reported an age estimate of 17.0-19 Ma for the Santacrucian SALMA based on a locality from northern Chile. If we accept the proposal of Croft *et al.* (2004, 2007) this would extend the Santacrucian SALMA to be between 16-19 Ma. The latter indicates that the 'Pinturan' and Santacrucian SALMA at least partially overlap. The recognition of a 'Pinturan' SALMA as a distinct biochronologic unit is still controversial (Kramarz *et al.*, 2010), and for that reason at this time we do not yet consider the 'Pinturan' SALMA to be a distinct and recognizable SALMA unit, and probably represents only a local fauna within the Santacrucian.

None of the terrestrial mammalian fauna from Cerro La Cruz recovered so far allows us to assign the fauna unequivocally to either the 'Pinturan' or Santacrucian SALMA as defined by Marshall et al. (1983). Neither the litoptern nor astrapothere specimens recovered so far can be identified to genus, let alone species, so they cannot be used to refine our understanding of where the fauna fits biostratigraphically. The known records of orophodonts are from the Deseadan SALMA of Argentina, while the pampathere Scirrotherium is known from Laventan SALMA, Colombia, the Late Miocene of Costa Rica (Laurito and Valerio, 2013), and the Huayquerian SALMA (Late Miocene ~9.0-6.8 Ma) of Argentina and Brazil (Dozo et al., 2008; Góis et al., 2013). So either the biostratigraphic range of the orophodonts is younger than previously known, and the stratigraphic range of Scirrotherium extends farther back in time. Neither pampatheres nor orophodonts are currently known from the classic Santacrucian or 'Pinturan' faunas.

So far the oldest known record of the fresh water turtle *Chelus* comes from the lower Miocene of Barzalosa Formation (Cundinamarca Department, Colombia), based on the xiphiplastron and costals referred to *Chelus colombiana* (Cadena *et al.*, 2008). Thus, the records of *Chelus* sp. from the Early Miocene of Cerro La Cruz, Castillo Formation confirms its early appearance in the fossil record of the north of South America (during the Early Miocene), and also represent the oldest record of this genus in Venezuela. The presence of the Serrasalmidae, *Colossoma* and *Mylossoma* during the Early Miocene of Cerro La Cruz represent one of the oldest records of these genera in the Neogene of South America (Lundberg *et al.*, 2010).

The levels from which the Cerro La Cruz mammals were recovered could be correlated with the C5Dr Magnetic Chron (Cande and Kent, 1995), which has an estimated age of 18 Ma (Burdigalian). Ortiz-Jaureguizar (1986) observed a direct link between the Colhuehuapian (early Burdigalian, ~20 Ma), and Santacrucian (late Burdigalian, ~17-16 Ma) SALMAs and grouped them into a single major evolutionary unit, called the Pansantacrucian faunal sub-cycle. This idea has been used by Bostelmann et al. (2013), for a succession of marine and continental strata from outcrops in southern Chile, with a rich assemblage of terrestrial vertebrate fossils, that are also biostratigraphically equivalent to a post-Colhuehuapian, pre-Santacrucian time interval, suggesting an age range of 19 to 17.8 Ma (Bostelmann et al., 2013), a situation similar to that of our current understanding of the biochronology at Cerro La Cruz based on the Strontium based ages presented her which places it between 17.21 (+0.41/-0.49) to 19.27 (+0.64/-0.53). At this time the finest time resolution possible for the locality is to consider this mammal assemblage as part of the Pansantacrucian mammalian subcycle.

Another locality in Venezuela that probably could be associated with Pansantacrucian mammalian sub-cycle includes the Quebrada Honda site from the Early Miocene of Chaguaramas Formation, eastern Venezuela, with a little known fauna composed by *Xenastrapotherium christi* (Stehlin, 1928), *Podocnemis geologorum* (Simpson, 1943), and indeterminate crocodilian remains (Stehlin, 1928; Wesselingh and Macsotay, 2006; Sánchez-Villagra *et al.*, 2010). In addition, the true age of the other two 'Early Miocene' Venezuelan localities that contain *Pseudoprepotherium venezuelanum* and *Boreostemma venezolensis* from Río Yuca and Quiamare Formation respectively remains unclear.

## 4.2. Paleoenvironments

In general, the Castillo Formation has been interpreted to have been deposited as an extensive coastal and marginal marine complex that includes a diverse range of paleoenvironments; terrestrial, fluvial, tidal, and brackish to open marine subtidal (Wheeler, 1960, 1963; Sánchez-Villagra *et al.*, 2000; Johnson *et al.*, 2009). However, the Cerro La Cruz sequence presents a wide variety of lithologies that have both lateral and vertical geometries that are consistent with Wheeler's (1960) ideas, and reflect

gradual changes in sedimentary environments being the result of eustatic changes.

The Cerro La Cruz sequence was deposited in the form of alternating packages of siliciclastic and carbonate sediments (Fig. 3). Its lower part is predominantly composed of siliciclastic sediments (mainly unfossiliferous mudstone along with a few interbeds of sandstones and conglomerate). Overlying, the sequence becomes predominately carbonate sediments represented by highly fossiliferous (mostly composed of marine invertebrates and fish otoliths), limestone strata (wackestone and packstone) with interbeddedby shale beds, and some sandstone, the mudstones (sometimes unfossiliferous and sometimes containing skeletal grains). This is followed by a siliciclastic-carbonate sequence predominantly represented by mudstones (sometimes fossiliferous) and sandstone interbedded with fossiliferous limestone beds (wackestone grainstone and packstone), with two levels containing 'continental or fresh water' vertebrate remains, leaves, and wood. The upper package is again primarily siliciclastic sediments, which is predominantly a shale sequence with intercalated beds of sandstone.

Bioturbation is common in the Cerro La Cruz section. Three main ichnofabrics are recognizing: Gyrolithes, Ophiomorpha and Thalassinoides, with Thalassinoides being conspicuous, and Gyrolithes more locally distributed. The paleoenvironmental distribution of Thalassinoides is quite varied, ranging from tidal flats, and both shoreline settings, to outer shelf facies, and even deep sea fan deposits (Ekdale et al., 1984; Myrow, 1995). Ophiomorpha are common in, but is by no means confined to or mainly indicative of beaches and near-shore sublittoral sands, as it also can occur in shoals, tidal flats, tidal stream point bars, and lagoon, bay, sound, and estuary floors, wherever the salinity and current energy are moderately high and the substrate consists mainly of sand (Frey et al., 1978). The Thalassinoides-Ophiomorpha ichofacies assemblage could be interpreted as having developed on subtidal substrates with well aerated agitated shallow water which are dominant in the most of the Cerro La Cruz section (Verde, 2002).

The *Gyrolithes* ichnofacies represents deep dwelling burrowers that are common in marginal-marine settings and its vertical helical morphology represents a specialized burrowing architecture that permits its makers to seek refuge from extreme salinity fluc-

tuations in brackish-water environments (Beynon and Pemberton, 1992; Buatois et al., 2005; Netto et al., 2007). The modern laomediid shrimp Axianassa australis sometimes produces spirals identical to Gyrolithes (Dworschak and Rodríguez, 1997; Felder, 2001), and five of the six known extant species of Axianassa inhabit intertidal zones, near mangrove areas (mud flats and channels) and only one species lives in deep-marine settings (Dworschak and Rodríguez, 1997; Netto, 2007). Toward the middle of the section analyzed, the presence of *Gyrolithes* combined with abundant wood remains (some associated with oysters in living positions) that we have observed in the middle and upper part of the unit reflects episodes of brackish-water probably lagoons near mangrove areas.

Transitions from siliciclastic to carbonate sedimentation such as that observed at Cerro La Cruz usually occurs mainly in near-coastal and inner shelf environments, and the changes from carbonate to non-carbonate deposition and vice versa are the result of variations in the supply of terrigenous clastic material and is determined by several factors like climatic, tectonic, or eustatic changes (Nichols, 2009).

The climate during the Late Oligocene to Early Miocene was a period characterized by relative higher global warmth possibly with higher atmospheric carbon dioxide (CO<sub>2</sub>) levels (Pagani et al., 1999; Paul et al., 2000). Wright and Miller (1992) have identified twelve significant  $\delta^{18}$ O increases in Oligocene and Early late Miocene benthic foraminiferal records (classified as Oi and Mi zones) that have been interpreted by several authors (Wright and Miller, 1992; Paul et al., 2000; Zachos et al., 2001) as indicative of the presence of large, transient Antarctic glaciations. The first and most prominent of these oxygen isotope excursions in the Miocene, Mi-1, occurred near the Oligocene/Miocene (O/M) boundary (Paul et al., 2000). Based on benthic foraminifera oxygen isotope records Wright and Miller (1992) also recognized that the maximum  $\delta^{18}$  values were associated with these increases at the base of two zones: Mi-1a and Mi-1b. More recently, Billups et al. (2002) dated the events Mi-1a at 21.69 Ma and Mi-1b at 18.0 Ma. Therefore the event Mi1b could be recorded in Cerro La Cruz sequence specifically in Facies C, and it is possible that the Mi-1b may be correlated with the presence of the continental and fresh water faunas present in the sequence.

The major part of the rocks of Falcón Basin was deposited during an extensional tectonic regimen dominated by tectonic collapse and graben formation, which was almost continuous since Late Eocene until Early Miocene. This extensional tectonic regimen was more pronounced during Late Eocene-Oligocene due to low-angle subduction in north of the region, between the current Caribbean Plate to the north and the Bonaire Block to the south (Baquero et al., 2009). The Cerro La Cruz sequence represents sedimentation of the south part of Falcón Basin deposited during the last episodes of the extensional tectonic regimen, with constant decline until the convergence of the Caribbean Plate from the north and the consequent tectonic inversion during the Middle Miocene, justifying the continuous decrease in sedimentation rate observed in the Unit B and Unit C compared with Unit A (Table 2, Fig. 3).

The Cerro La Cruz sequence includes a time period of at least 2 My. During this interval, according to several authors (Haq et al., 1987; Miller et al., 1998; Kominz et al., 1998; Miller et al., 2005; Browning et al., 2008; Kominz et al., 2008), the sea level started to regress between ~19-18 Ma, follow by a transgression starting about ~18 until ~17 Ma. This sequence could be correlated with the dynamics of the change in the facies present at Cerro La Cruz. Facies A and B represent the regressive phase (probably with some oscillations, Fig. 3), and facies C a trangressive phase followed by a later regression with eustatic variations. Most of the Cerro La Cruz faunas would be associated with the start of the transgression around 18 Ma ago, and to the Mi-1b event of Antarctic glaciations.

Generally the near-shore marine environments (subtidal shallow marine, lagoon, intertidal zone near mangroves) are dominant in the Cerro La Cruz sequence, but the mammal assemblage from the two levels from the middle part of the section indicate the presence of continental environments, although for a short duration. Ecologically, the Orophodontidae, Astrapotheriinae and Litopterna represent forest taxa, while members of the Pampatheriidae are considered grassland inhabitants. Other taxa with fresh water affinities present at Cerro La Cruz are the Characiform (Serraselmidae) fish Colossomma and Mylossoma with multicuspid molariform teeth, which facilitate crushing fruits and hard-coated seeds (Correa et al., 2007). According to Dahdul (2004) Colossoma macropomum is present in Cerro La Cruz fauna.

This is an extant species that does occur in muddy-or black-water rivers but is limited to clear-water, shield-draining rivers in their lower reaches, often below large waterfalls or cataracts (Araújo-Lima and Goulding, 1997). Both *Colossoma* and *Mylossoma* have a diet based on vegetable material like fruits and hard-coated seeds, which it takes from the water surface of rivers, under the forest cover (Useche *et al.*, 1993). The extant species of the turtle *Chelus fimbriata* is a highly aquatic, lowland tropical species, found equally in rivers in both forested areas and savannah habitats in low tropical regions of Venezuela (Pritchard, 2008). Living species of *Colossomma*, *Mylossoma* and the turtle *Chelus* could be found in modern Orinoco and Amazon Basin.

The presence of fresh water and continental faunal assemblage suggests a predominance of forest with probably some savanna areas, and some rivers in the Cerro La Cruz, which agree with Strömberg *et al.* (2013), who using a high-resolution record of plant silica (phytoliths) from Patagonia described a dominance of forest areas (at least 70%) with minor open-habitat grasses (about 5 to 30%) in South America during the Early Miocene.

# 4.3. Implications of the 'Orinoco River shifting' hypothesis

The dynamic geologic history of South America should thus be very relevant for understanding the origins of the present diversity in the Amazonia (Hoorn et al., 2010). During the Cenozoic an ancient river-dominated landscape usually called the 'Proto-Orinoco' or 'Proto-Amazon' was present in the northern of South America, collecting most of the Colombian, Ecuadorian and Peruvian Amazon tributaries and flowing mostly towards the Caribbean Sea (Rod, 1981; Hoorn et al., 1995; Díaz de Gamero, 1996; Lundberg et al., 1998; Hooghiemstra et al., 2006; Wesselingh and Salo, 2006; Hoorn and Wesselingh, 2010; Shephard et al., 2010; Hoorn et al., 2010; Mora et al., 2010; Aguilera et al., 2013). Based on the fluvial and deltaic sediments of Paleocene age of the Orocue Group and the Marcelina Formation, the Eocene sediments of the Mirador Formation from the Venezuelan Llanos Basin, and the vast deltaic sedimentation of the Misoa and Carbonera Formations (Eocene) in the Maracaibo Basin, the existence of a large river system flowing in a general south-north direction during Paleocene to

Eocene in the Maracaibo Basin is accepted (Kasper and Larue, 1986; Díaz de Gamero, 1996).

Later, during Oligocene or Miocene times the course of this major river system changed to a clockwise direction until the present day position (Rod, 1981; Díaz de Gamero, 1996). Commonly, this change has been attributed to the Andean uplift (Hoorn *et al.*, 1995; Díaz de Gamero, 1996; Hoorn *et al.*, 2010), or driven by mantle convection (Shephard *et al.*, 2010). But the timing of this change is still unclear.

Different hypothesis exist regarding the evolution of the north of the 'Proto-Orinoco' River during Oligocene/Miocene to its present-day position (Guzmán and Fisher, 2006). The first hypothesis considered that until the Early Miocene, most of the area of the contemporary western Amazon drained northward to a delta located in the area of the modern Maracaibo Basin (Hoorn et al., 1995; Lundberg, 1998; Toro and Steel, 2002; Albert et al., 2006). However, the initial uplift and exhumation of the Colombian Eastern Cordillera during Oligocene and the paleogeographic reconstructions based on structural, stratigraphic, and sedimentological data from Maracaibo Basin and other Venezuela and Colombia basins shows no evidence for deposition of thick and extensive deltaic sedimentary deposits that would be expected with the outflow of a major river system similar to modern Orinoco or Amazon rivers (Villamil, 1999; Parra et al., 2005; Guzmán and Fisher, 2006). The recent Orinoco basin drains 75% of Venezuelan territory and most of the eastern lowlands of Colombia, covering an approximate area of 1×106 km<sup>2</sup> (Meade, 1994; Méndez, 2005). With approximately 30.000 km<sup>2</sup> the Orinoco delta is one of the largest deltas in America (Méndez, 2005). The recent Orinoco River ranks third in the world in terms of water discharge and about tenth in sediment discharge, with an average of suspended-sediment load of about 150×106 tons (Meade, 1994). These ideas discard a paleo-Maracaibo outlet for most of the majors rivers flowing north during the Early Miocene (Johnson et al., 2009; Mora et al., 2010).

A second hypothesis proposed that during the Oligocene-Miocene boundary the major river system ('Proto-Orinoco') drained into the Falcón Basin northwestern of Venezuela (Rod, 1981; Díaz de Gamero, 1996; Aguilera *et al.*, 2013). It is due to some geological evidence from the Falcón Basin, besides the biogeographical relationships of the rich fresh-water

vertebrate fossil fauna found in the northwestern Falcón (Socorro, Urumaco, Codore and Tío Gregorio Formations) to the recent Orinoco River system (Díaz de Gamero, 1996). Among others, Díaz de Gamero (1996) proposed that the clastics sediments present in the Castillo Formation, western margin of Falcón Basin, probably represent the fluvio-deltaics facies of the 'Proto-Orinoco' River. Nonetheless, Díaz de Gamero (1996) notes that the Castillo Formation was poorly studied, and this author, following Wheeler (1960) reported the presence of coal seams, and interpreted the formation as fluvial deltaic and coastal marine. But Wheeler (1960, 1963) described only the continental to brackish shales (along the basin edge) that contain a few thin coal seams, and in more than 360 m of stratigraphic section at Cerro La Cruz outcrops we did not observe any coals seam level. Furthermore, based in the presence of the Serrasalmidae fishes Colossoma and Mylossoma found in Cerro La Cruz, Dahdul (2004), Aguilera and Rodríguez de Aguilera (2004a), and Sánchez-Villagra et al. (2004) suggest that records of 'Proto-Orinoco' shift is contained in the Miocene sections of Cerro La Cruz, Castillo Formation. Also Aguilera et al. (2013) proposed the presence of a major river system in the western margin of the Falcón Basin during the earliest Miocene, with clear faunal affinities to the Amazon and Orinoco, probably referring to the Cerro La Cruz fauna.

In contrast with the last two hypotheses, the geological settings and the paleontological affinities of the vertebrate fauna so far recovered from Cerro La Cruz (Castillo Formation, Early Miocene, Falcón Basin) do not provide any conclusive proxy to indicate the presence of a major river system that could be interpreted as the 'Proto-Orinoco'. Nevertheless, the occurrence of some freshwater taxa such as Colossoma, Mylossoma, and Chelus from Cerro La Cruz suggests the presence of local fluvial environments associated to these fresh water faunas, and the fauna so far recovered from Cerro La Cruz shows Orinoco-Amazon affinities. This is based on the presence of extant species of Colossoma, Mylossoma, and Chelus in the recent Orinoco Basin, although some of these genera can be found in other hydrographic basins such as Maracaibo Basin, where Mylossoma acanthogaster, and Chelus fimbriata have been found.

Thus we consider unlikely a direct connection between the Cerro La Cruz sections with a major river system during the Early Miocene. The inferred direction of the flow of this local fluvial environment (river) was to the north. The limited clastic terrigenous influx and the immature sandstone observed in the analyzed section suggest this fluvial environment probably had a low water volume and a short path. We consider more probable that Cerro La Cruz was part of the same hydrographic basin that gave rise to the modern Orinoco River before Miocene, but being isolated at least since the Early Miocene.

Observations from Cerro La Cruz support a third hypothesis, which indicated that the drainage axis of the major river system ('Proto-Orinoco' River) had been diverted to the northeast since the Late Oligocene (Villamil, 1999). Furthermore, Guzman and Fisher (2006) propose that the Maracaibo Basin has been isolated from extrabasinal drainage systems since the Late Oligocene, and sediments derived from the surrounding highlands were either deposited in the basin or delivered into the neighboring Falcón Basin through a narrow marine passage. So the Oligocene-Miocene intrabasinal drainage systems in the Maracaibo and Falcón Basins support that the course of the 'Proto-Orinoco' River must have been permanently shifted away from the Maracaibo Basin since the Oligocene or earlier. This third hypothesis was also supported by the presence of the bivalve Pachydon hettneri in the Chaguaramas Formation (Late Oligocene/Early Miocene, with outcrops in Guárico State, Eastern of Venezuela), Perú and Colombia, implying that during the Burdigalian-Langhian a lowland aquatic biogeographic connection existed between the Amazon region and Venezuela through the East Andean foreland basins (Wesselingh and Macsotay, 2006). In addition, the sandstones and gravels of the Chaguaramas Formation contain a heavy mineral suite characteristic of the Colombian Central Cordillera, and the clay mineralogy of the silt component of this formation indicates both a Colombian Central Cordilleran and a Venezuelan coastal range origin (Vivas and Macsotay, 1995; Wesselingh and Macsotay, 2006). Consequently, the sedimentological and paleontological records suggest that the Chaguaramas Formation could be considered as relict of 'Proto-Orinoco' delta (Vivas and Macsotay, 1995; Pindell et al., 1998; Wesselingh and Macsotay, 2006), discarding the possible draining of a major river system ('Proto-Orinoco') to the Maracaibo or Falcón Basin during the Early Miocene.

#### 5. Conclusions

The recorded fauna of Cerro La Cruz includes 40 taxa: nine Chondrichthyes, 13 Actinopterygii, four Testudines, four Cetartiodactyla, three Crocodilia, one Sirenia, three Xenarthra, two Notoungulata and one Litoptern. The last six confirm the presence of the earliest and most diverse terrestrial mammal assemblage in Venezuela. The geologic analysis of the sediments present at Cerro La Cruz allows us to interpret the upper part of the Castillo Formation, with outcrops in the southwestern Sierra de la Baragua, Lara state, as a mainly near-shore marine complex with local brackish-water probably lagoon near mangrove areas, subtidal shallow marine environments, shelf lagoon with low energy water circulation, and shallow water with open circulation close to wave base, associated to regressive and trangressive phases probably due to Antarctic glaciations, during Early Miocene (87Sr/86Sr ages between 17.21-19.27 Ma). Nevertheless, at least two different continental episodes, that contains an Early Miocene mammalian assemblage were found. The number of terrestrial mammalian taxa discovered to date, does not permit assignment to a specific SALMA, although does fit within the earlier part of the broader defined Pansantacrucian mammalian sub-cycle. The continental vertebrate assemblage is interpreted as indicative of an ecosystem consisting of a mosaic that includes dominant tropical humid forest and some lowland savanna.

Our interpretation of the geology at Cerro La Cruz shows no conclusive evidence for the presence of a big river crossing over that zone during the Early Miocene, and does not support the hypothesis of the possible draining of a 'Proto-Orinoco' river into Maracaibo or Falcón Basin during Early Miocene. The recorded vertebrate fauna indicate some similarities with other northern South America fossil localities with Orinoco-Amazon affinities, the Middle-Late Miocene fauna of La Venta (Colombia) and Urumaco (Venezuela), nevertheless futures analyses will be needed corroborate this hypothesis.

#### Acknowledgments

We wish to thank to Instituto del Patrimonio Cultural (IPC), Venezuela, for permission for this project. We thank to Sr. P. Gómez<sup>†</sup>, Ing. M.E. Mendoza, Lic. M. López and Dr. F. Urbani with their help and camaraderie during the field trips. This Project is supported by a grant from TOTAL Venezuela and the Instituto Venezolano de

Investigaciones Científicas (IVIC) under the project 822 and 1096 to ADR. We also thank to T. Hernández from Istituto de Geofísica, UNAM, for all his help in processing the samples to calculate the age of strontium.

#### References

- Aguilera, O.; Lundberg, J. 2010. Venezuelan caribbean and orinocoan neogene fish. *In* Urumaco and Venezuelan Paleontology, The Fossil Record of the Northern Neotropics (Sánchez-Villagra, M.; Aguilera, O.; Carlini, A.A.; editors). Indiana Press University: 129-152.
- Aguilera, O.; Rodríguez de Aguilera, D. 2004a. Amphi-American Neogene sea catfishes (Siluriformes, Ariidae) from Northern South America. *In Fossils of the Miocene Castillo Formation*, Venezuela: contributions on neotropical palaeontology (Sánchez-Villagra, M.R.; Clack, J.A.; editors). Special Papers in Palaeontology 71: 29-48.
- Aguilera, O.; Rodríguez de Aguilera, D. 2004b. New Miocene otolith-based Sciemid species (Pisces, Perciformes) from Venezuela. *In Fossils of the Miocene* Castillo Formation, Venezuela: contributions on neotropical palaeontology (Sánchez-Villagra, M.R.; Clack, J.A.; editors). Special Papers in Palaeontology 71: 49-59.
- Aguilera, O.; Rodríguez de Aguilera, D.; Vega, F.J.; Sánchez-Villagra, M. 2010. Mesozoic and Cenozoic decapod crustaceans from Venezuela and related trace-fossil assemblages. *In* Urumaco and Venezuelan Paleontology, The Fossil Record of the Northern Neotropics (Sánchez-Villagra, M.; Aguilera, O.; Carlini, A.A.; editors). Indiana Press University: 103-129.
- Aguilera, O.; Moraes-Santos, H.; Costa, S.; Ohe, F.; Jaramillo, C.; Nogueira, A. 2013. Ariid sea catfishes from the coeval Pirabas (Northeastern Brazil), Cantaure, Castillo (Northwestern Venezuela), and Castilletes (North Colombia) formations (Early Miocene), with description of three new species. Swiss Journal of Palaeontology 132 (1): 45-68.
- Albert, J.S.; Lovejoy, N.R.; Crampton, W.G.R. 2006. Miocene tectonism and the separation of cis- and trans-Andean river basins: Evidence from Neotropical fishes. Journal of South American Earth Science 21: 14-27.
- Araujo-Lima, C.; Goulding, M. 1997. So Fruitful a Fish, Ecology, Conservation and Aquaculture of the Amazon's Tambaqui. Columbia University Press: 191 p. New York.
- Audemard, F.A. 1993. Neotectonique, Sismotectonique et Ale'a Sismique du Nord-ouest du Venezuela (systeme

- de failles d'Oca-Ancon). Ph.D. Thesis (Unpublished), Universite Montpellier II: 369 p.
- Audemard, F.A. 2001. Quaternary tectonics and present stress tensor of the inverted northern Falcon Basin, northwestern Venezuela. Journal of Structural Geology 23: 431-453.
- Baquero, M.; Acosta, J.; Kassabji, E.; Zamora, J.; Sousa, J.; Rodríguez, J.; Grobas, J.; Melo, L.; Shneider, F. 2009. Polyphase development of the Falcón basin in Northwestern Venezuela, implications for oil generation. *In* The origin and evolution of the Caribbean plate (James, K.H.; Lorente, M.A.; Pindell, J.L.; editors). Geological Society of London Special Publications 328: 587-612.
- Beynon, B.M.; Pemberton, S.G. 1992. Icnological signature of a brackish water deposit, an example from the Lower Cretaceous Grand Rapids Formation, Cold Lake Oil Sands area, Alberta. *In Applications of Ichnology to petroleum exploration (Pemberton S.G.*; editor). Society of Sedimentary Geologists: 199-221.
- Billups, K.J.; Channell, E.T.; Zachos, J. 2002. Late Oligocene to Early Miocene geochronology and paleoceanography from the subantarctic South Atlantic. Paleoceanography 17 (1): 4-11.
- Bond, M.; Gelfo, J.N. 2010. The South American native ungulates of the Urumaco Formation. *In* Urumaco and Venezuelan Paleontology, The Fossil Record of the Northern Neotropics (Sánchez-Villagra, M.; Aguilera, O.; Carlini, A.A.; editors). Indiana Press University: 256-268.
- Bostelmann, J.E.; Le Roux, J.P.; Vásquez, A.; Gutiérrez, N.; Oyarzún, J.; Carreño, C.; Torres, T.; Otero, R.; Llanos, A.; Fanning, M.; Hervé, F. 2013. Burdigalian deposits of the Santa Cruz Formation in the Sierra Baguales, Austral (Magallanes) Basin: Age, depositional environment and vertebrate fossils. Andean Geology 40 (3): 458-489.
- Brochu, C.A.; Rincón, A.D. 2004. A Gavialoid Crocodylian from Lower Miocene of Venezuela. *In* Fossils of the Miocene Castillo Formation, Venezuela: contributions on neotropical palaeontology (Sánchez-Villagra, M.R.; Clack, J.A.; editors). Special Papers in Palaeontology 71: 61-79.
- Browning, J.V.; Miller, K.G.; Sugarman, P.J.; Kominz, M.A.; McLaughlin, P.P.; Kulpecz, A.A. 2008. 100 Myr record of sequences, sedimentary facies and sealevel change from Ocean Drilling Program onshore coreholes, U.S. Mid-Atlantic coastal plain. Basin Research 20: 227-248.

- Buatois, L.A.; Gingras, M.K.; Maceachern, J.; Mángano, M.G.; Zonneveld, J.P.; Pemberton, S.G.; Netto, R.G.; Martin, A.J. 2005. Colonization of brackish-water systems through time, evidence from the trace-fossil record. Palaios 20: 321-347.
- Cadena, E.; Jaramillo, C.; Paramo, M.E. 2008. New material of *Chelus colombiana* (Testudines; Pleurodira) from the lower Miocene of Colombia. Journal of Vertebrate Paleontology 28 (4): 1206-1212.
- Cande, S.C.; Kent, D.V. 1995. Revised calibration of the geomagnetic polarity time scale for the late Cretaceous and Cenozoic. Journal of Geophysical Research 100: 6093-6095.
- Cati, F.; Colalongo, M.L.; Crescenti, U.; D'Onofrio, S.; Follador, U.; Pirini, R.; Pomesano, C.; Salvarorini, G.; Sartorini, S.; Premoli Silva, I.; Wezel, C.F.; Bertolino, V.; Bizon, G.; Bolli, H.M.; Borsetti, C.A.M.; Dondi, F.L.H.; Jenkins, D.G.; Perconing, E.; Sampo, M.; Sprovieri, R. 1968. Biostratigrafia del Neogeno mediterráneo basata sui foraminiferi planctonici. Bollettino della Societa Geologica Italiana 87: 491-503.
- Collins, R.L. 1934. Venezuela Tertiary mammals. Johns Hopkins University Studies in Geology 2: 235-244.
- Correa, S.; Kirk, W.; López-Fernández, H.; Galetti, M. 2007. Evolutionary perspectives on seed consumption and dispersal by dishes. BioScience 57 (9): 748-756.
- Croft, D.A.; Flynn, J.J.; Wyss, A.R. 2004. Notoungulata and Litopterna of the Early Miocene Chucal fauna, northern Chile. Fieldiana Geology 50: 1-52.
- Croft, D.A.; Flynn, J.J.; Wyss, A.R. 2007. A new basal glyptodontid and other Xenarthra of the Early Miocene Chucal fauna, northern Chile. Journal of Vertebrate Paleontology 27: 781-797.
- Dahdul, W.M. 2004. Fossil serrasalmine fishes (Teleostei, Characiformes) from the Lower Miocene of Northwestern Venezuela. *In* Fossils of the Miocene Castillo Formation, Venezuela: contributions on neotropical palaeontology (Sánchez-Villagra, M.R.; Clack, J.A.; editors). Special Papers in Palaeontology 71: 23-28.
- Díaz de Gamero, M.L. 1996. The changing course of the Orinoco River during the Neogene, a review. Palaeogeography, Palaeoclimatology, Palaeocology 123: 385-402.
- Dott, R.H. 1964. Wacke, graywacke and matrix-what approach to inmature sandstone classification. Journal of Sedimentary Petrology 34: 625-632.
- Dozo, M.T.; Bouza, P.; Tambussi, C.; Massaferro, G. 2008. Vertebrados continentales de Edad Huayqueriense (Mioceno Tardío) en Península Valdés (Chubut, Argentina): Implicaciones biocronológicas, paleoam-

- bientales y paleobiogeográficas. *In* Congreso Latinoamericano de Paleontología, No. 3: 83 p. Neuquén.
- Droser, M.L.; Bottjer, D.J. 1986. A semiquantitative field classification of ichnofabric. Journal of Sedimentary Research 56: 558-559.
- Dunham, R.J. 1962. Classification of carbonate rocks according to depositional texture. *In Classification of carbonate rocks* (Ham, W.E.; editors). American Association of Petroleum Geologists, Memoir: 108-121.
- Dunn, R.; Madden, R.; Kohn, M.; Schmitz, M.; Strömberg, C.; Carlini, A.; Ré, G.; Crowley, J. 2012. A new chronology for middle Eocene-Early Miocene South American Land Mammal Ages. Geological Society of America, Bulletin 125 (3-4): 539-555.
- Dworschak, P.C.; Rodríguez, S. 1997. A modern analogue for the trace fossil *Gyrolithes*, Burrows of the thalassinidean shrimp *Axianassa australis*. Lethaia 30: 41-52.
- Ekdale, A.A.; Bromley, R.G.; Pemberton, S.G. 1984. Ichnology, The use of trace fossils in sedimentology and stratigraphy. Society of Economic Paleontologists and Mineralogists, Short Course 15: 317 p.
- Fasola, A.; Giffunni, G.; Crespo, S.; Paredes, I.; Uribe, A. 1985. Estudios bioestratigráficos del intervalo Cretáceo superior (Maestrichtiense) a Mioceno inferior en el norte del estado Guárico, Venezuela. *In* Congreso Geológico Venezolano, No. 6, Actas 1: 588-645. Caracas.
- Felder, D.L. 2001. Diversity and ecological significance of deep burrowing macrocrustaceans in coastal tropical waters of the Americas (Decapoda, Thalassinidea). Interciencia 26 (10): 440-449.
- Flügel, E. 2004. Microfacies of Carbonate Rocks, Analysis, Interpretation, and Application. Springer: 996 p. Berlín.
- Flynn, J.J.; Swisher, C.C. 1995. South American Land Mammal Ages: Correlation to global geochronologies. *In* Geochronology, Time Scales and Global Stratigraphic Correlation (Berggren, W.A.; editor). Society for Sedimentary Geology, Special Publication 54: 317-333.
- Frey, R.; Howard, J.; Pryor, W. 1978. Ophiomorpha, Its morphologic, taxonomic, and environmental significance. Palaeogeography, Palaeoclimatology, Palaeoecology 23: 199-229.
- Góis, F.L.; Scillato-Yané, G.J.; Carlini, A.A.; Guilherme, E. 2013. A new species of *Scirrotherium* Edmund and Theodor, 1997 (Xenarthra, Cingulata, Pampatheriidae) from the late Miocene of South America. Alcheringa 37: 177-188.
- Graham, I.J.; Morgans, H.E.G.; Waghorn, D.B.; Trotter, J.A.; Whitford, D.J. 2000. Strontium isotope stratigraphy of the Oligocene-Miocene Otekaike limestone (Trig Z section) in southern New Zealand, age of the

- Duntroonian/Waitakian Stage boundary. New Zealand Journal of Geology and Geophysics 43 (3): 335-347.
- Guzman, J.; Fisher, W. 2006. Early and middle Miocene depositional history of the Maracaibo Basin, western Venezuela. American Association of Petroleum Geologists, Bulletin 90 (4): 625-655.
- Haq, B.U.; Hardenbol, J.; Vail, P.R. 1987. Chronology of fluctuating sea levels since the Triassic (250 million years ago to present). Science 235: 1156-1167.
- Hoffstetter, R. 1969. Remarques sur la phylogénie et la classification des Edentés Xénarthres (Mammifères) actuels et fossils. Bulletin du Muséum National d'Histoire Naturelle 41: 91-103. Paris.
- Hooghiemstra, H.; Wijninga, V.; Cleef, A. 2006. The paleobotanical record of Colombia, implications for biogeography and biodiversity. Annals of the Missouri Botanical Garden 93 (2): 297-325.
- Hoorn, C.; Wesselingh, H. 2010. Amazonia, Landscape and Species Evolution, a Look into the Past (Hoorn, C.; Wesselingh, H.; editors). Wiley-Blackwell: 482 p. Oxford.
- Hoorn, C.; Guerrero, J.; Sarmiento, G.A.; Lorente, M.A. 1995. Andean tectonics as a cause for changing drainage patterns in Miocene northern South America. Geology 23: 237-240.
- Hoorn, C.; Wesselingh, F.P.; ter Steege, H.; Bermúdez, M.A.; Mora, A.; Sevink, J.; Sanmartín, I.; Sánchez-Meseguer, A.; Anderson, C.L.; Figueiredo, J.P.; Jaramillo, C.; Riff, D.; Negri, F.R.; Hooghiemstra, H.; Lundberg, J.; Stadler, T.; Särkinen, T.; Antonelli, A. 2010. Amazonia Through Time: Andean Uplift, Climate Change, Landscape Evolution, and Biodiversity. Science 330: 927-931.
- Howarth, R.J.; McArthur, J.M. 1997. Statistics for strontium isotope stratigraphy, a robust lowess fit to the marine Sr-isotope curve for 0 to 206 Ma, with look-up table for derivation of numeric age. Journal of Geology 105: 441-456.
- Johnson, K.; Sánchez-Villagra, M.; Aguilera, O.A. 2009. The Oligocene-Miocene Transition on Coral Reefs in the Falcón Basin (NW Venezuela). Palaios 24 (1): 59-69.
- Kasper, D.; Larue, D. 1986. Paleogeographic and tectonic implications of quartzose sandstones of Barbados. Tectonics 5 (6): 837-854.
- Kominz, M.A.; Miller, K.G.; Browning, J.V. 1998. Long-term and short-term global Cenozoic sea-level estimates. Geology 26: 311-314.
- Kominz, M.A.; Browning, J.V.; Miller, K.G.; Sugarman, P.J.; Misintseva, S.; Scotese, C.R. 2008. Late Creta-

- ceous to Miocene sea-level estimates from the New Jersey and Delaware coastal plain coreholes, an error analysis. Basin Research 20: 211-226.
- Kramarz, A.; Vucetich, M.G.; Carlini, A.; Ciancio, M.; Abello, M.A.; Deschamps, C.; Gelfo, J. 2010. A new mammal fauna at the top of the sequence at Gran Barranca, its biochronological meaning. *In* The Paleontology of Gran Barranca, Evolution and Environmental Change through the Middle Cenozoic of Patagonia (Madden, R.H.; Carlini, A.A.; Vucetich, M.G.; Kay, R.F.; editors). Cambridge University Press: 264-277.
- Laurito, C.A.; Valerio, A.L. 2013. Scirrotherium antelucanus, una nueva especie de Pampatheriidae (Mammalia, Xenarthra, Cingulata) del Mioceno Superior de Costa Rica, América Central. Revista de Geología de América Central 49: 45-62.
- López, V.M.; Brineman Jr., J.H. 1943. Estudio geológico y minero del yacimiento de mercurio de San Jacinto, estado Lara. Ministerio de Fomento, Revista de Fomento 5 (50): 29-61. Caracas.
- Lorente, M.A. 1987. Palynology and Palynofacies of the Upper Tertiary In Venezuela. Dissertationes Botanicae 99: 222 p. Berlín.
- Lundberg, J.G.; Marshall, L.G.; Guerrero, J.; Horton, B.; Malabarba, M.C.S.L.; Wesselingh, F. 1998. The stage for Neotropical fish diversification, a history of tropical South American rivers. *In* Phylogeny and Classification of Neotropical Fishes (Reis, R.E.; Vari, R.P.; Lucena, Z.M.; Lucena, C.A.S.; editors). Edipucrs: 13-48. Porto Alegre.
- Lundberg, J.G.; Sabaj-Pérez, M.H.; Dahdul, W.M.; Aguilera, O.A. 2010. The Amazonian Neogene fish fauna. *In* Amazonia, Landscape and Species Evolution, a Look into the Past (Hoorn, C.; Wesselingh, H.; editors). Wiley-Blackwell: 482 p. Oxford.
- Mactosay, O.; Ferry, S.; Fierro, I. 1989. Las formaciones Ranchería y Trujillo: evidencias sedimentológicas de su relación a la tectónica de obducción. *In* Congreso Geológico de Venezuela, No. 7, 1: 467-494. Barquisimeto.
- Macsotay, O.; Vivas, V.; Werhrmann, M.; Hartenberger, J.L.; Chachati, B. 1995. Tectono-sedimetary molasse cycles along northern Venezuela and Trinidad. *In* Transactions Geological Conference of the Geological Society, No. 3, of Trinidad and Tobago and the Caribbean Geological Conference, No. 14 (Ali, W.; Oli, A.; Young On, V.; editors). Held in Port of Spain, Trinidad: Geological Society of Trinidad and Tobago 2: 584-593.

- Marshall, L.G.; Hoffstetter, R.; Pascual, R. 1983. Mammals and stratigraphy, geochronology of the continental mammal-bearing tertiary of South America. Palaeovertebrata Mémoire Extraordinaire: 93 p.
- Marshall, L.G.; Drake, R.E.; Curtis, G.H.; Butler, R.F.; Flanagan, K.M.; Naeser, C.W. 1986. Geochronology of Type Santacrucian (Middle Tertiary) Land Mammal Age, Patagonia, Argentina. Journal of Geology 94 (4): 449-457.
- Martínez, G.; Valletta, G. 2008. Petrografia de las facies gruesas de la Formación Matatere y otras unidades del centro-occidente de Venezuela. Geos, Universidad Central de Venezuela 40 (201): 65-66. Caracas.
- Meade, R.H. 1994. Suspended sediments of the modern Amazon and Orinoco rivers. Quaternary International 21: 29-39.
- Méndez, J. 2005. El Delta Del Orinoco. *In* Frente Atlántico Venezolano, Investigaciones Geoambientales: Ciencias de la Tierra (Gómez, M.G.; Capaldo, M.; Yanes, C.; editors). Fondo Editorial Fundambiente 2: 12-24. Caracas.
- Miller, K.G.; Mountain, G.S.; Browning, J.V.; Kominz, M.; Sugarman, P.J.; Christie-Blick, N.; Katz, M.E.; Wright, J.D. 1998. Cenozoic global sea level, sequences, and the New Jersey transect, Results from coastal plain and continental slope drilling. Reviews of Geophysics 36: 569-601.
- Miller, K.G.; Kominz, M.; Browning, J.V.; Wright, J.; Mountain, G.; Katz, M.; Sugarman, P.J.; Cramer, B.; Christie-Blick, N.; Pekar, S. 2005. The Phanerozoic Record of Global Sea-Level Change. Science 310: 1293-298.
- Mora, A.; Baby, P.; Roddaz, M.; Parra, P.; Brusset, S.; Hermoza, W.; Espurt, N. 2010. Tectonic history of the Andes and sub-Andean zones: implications for the development of the Amazon drainage basin. *In* Amazonia, Landscape and Species Evolution, a Look into the Past (Hoorn, C.; Wesselingh, H.; editors). Wiley-Blackwell: 482 p. Oxford.
- Muessig, K.W. 1984. Structure and Cenozoic tectonics of the Falcón Basin, Venezuela, and adjacent areas. *In* The Caribbean-South American Plate Boundary and Regional Tectonics (Bonini, W.E.; Hargraves, R.B.; Shagam, R.; editors). Geological Society of America, Memoirs 162: 217-230.
- Myrow, P. 1995. *Thalassinoides* and the Enigma of Early Paleozoic Open-Framework Burrow Systems. Palaios 10 (1): 58-74.
- Netto, R.G.; Buatois, L.A.; Gringas, M.K.; Mángano, M.G.; Ballisteri, P. 2007. *Gyrolithes* as a multipurpose

- burrow, an ethologic approach. Revista Brasileira de Paleontologia 13: 61-69.
- Nichols, G. 2009. Sedimentology and Stratigraphy. Wiley-Blackwell Publication, second edition: 419 p.
- O'Leary, M.A. 2004. A fragmentary odontocete cranium from the Lower Miocene of Venezuela. *In* Fossils of the Miocene Castillo Formation, Venezuela: contributions on neotropical palaeontology (Sánchez-Villagra, M.R.; Clack, J.A.; editors). Special Papers in Palaeontology 71: 99-104.
- Ortiz-Jaureguizar, E. 1986. Evolución de las comunidades de mamíferos cenozoicos sudamericanos: un estudio basado en técnicas de análisis multivariado. *In* Congreso Argentino de Paleontología y Bioestratigrafía, No. 4, Actas 4: 191-207. Mendoza.
- Oslick, J.S.; Miller, K.G.; Feigenson, M.D. 1994. Oligocene Miocene strontium isotopes, Stratigraphic revisions and correlations to an inferred glacioeustatic record. Paleoceanography 9 (3): 427-443.
- Pagani, M.; Arthur, M.A.; Freeman, K.H. 1999. Miocene evolution of atmospheric carbon dioxide. Paleoceanography 14 (3): 273-292.
- Parra, M.; Mora, A.; Jaramillo, C.; Veloza, G.; Strecker, R. 2005. New stratigraphic data on the initiation of mountain building at the Eastern Front of the Colombian Eastern Cordillera. *In* International Symposium on Andean Geodynamics, Extended Abstracts: 567-571. Barcelona.
- Patterson, B. 1977. A primitive pyrothere (Mammalia, Notoungulata) from the early Tertiary of Northwestern Venezuela. Fieldiana Geology 33: 397-422.
- Paul, H.; Zachos, J.; Flower, B.; Tripati, A. 2000. Orbitally induced climate and geochemical variability across the Oligocene/Miocene boundary. Paleoceanography 15 (5): 471-485.
- Pindell, J.L.; Higgs, R.; Dewey, J.F. 1998. Cenozoic palinspastic reconstruction, paleogeographic evolution and hydrocarbon setting of the northern margin of South America. Society for Sedimentary Geology (SEPM), Special Publication 58: 45-85.
- Pritchard, P.C.H. 2008. *Chelus fimbriata* (Schneider 1783) matamata turtle. *In* Conservation Biology of Freshwater Turtles and Tortoises: A Compilation Project of the IUCN/SSC Tortoise and Freshwater Turtle Specialist Group (Rhodin, A.G.J.; Pritchard, P.C.H.; van Dijk, P.P.; Saumure, R.A.; Buhlmann, K.A.; Iverson, J.B.; editors). Chelonian Research Monographs 5: 1-10.
- Ré, G.H.; Bellosi, E.S.; Heizler, M.; Vilas, J.F.; Madden, R.H.; Carlini, A.A.; Kay, R.F.; Vucetich, M.G. 2010. A geochronology for the Sarmiento Formation at Gran

- Barranca. *In* The Paleontology of Gran Barranca: Evolution and Environmental Change trough the Middle Cenozoic of Patagonia (Madden, R.H.; Carlini, A.A.; Vucetich, M.G.; Kay, R.F.; editors). Cambridge University Press: 46-58.
- Rincón, A.D.; Solórzano, A.; Mendoza, M. 2010a. El registro de vertebrados del Mioceno temprano de la Formación Castillo, estado Lara, Venezuela. *In* Simposio, No. 40, Aniversario del Centro de Ecología, Instituto Venezolano de Investigaciones Científicas, Resúmenes: p. 120.
- Rincón, A.D.; Solórzano, A.; McDonald, H.G. 2010b. New mammal records from the Early Miocene Castillo Formation of Venezuela. *In* Anniversary, No. 70, Meeting Society of Vertebrate Paleontology Pittsburgh, Special Issue 28 (3): 151a. Pennsylvania.
- Rod, E. 1981. Note on the shifting course of the ancient Río Orinoco from Late Cretaceous to Oligocene time. Geos 26: 54-56.
- Sánchez-Villagra, M.R.; Burnham, R.J.; Cambell, D.C.; Feldmann, R.M.; Gaffney, E.S.; Kay, R.F.; Lozsán, R.; Purdy, R.; Thewissen, J.G.M. 2000. A new near-shore marine fauna and flora from the early Neogene of northwestern Venezuela. Journal of Paleontology 74 (5): 957-968.
- Sánchez-Villagra, M.; Gasparini, Z.; Lozsán, R.; Moody, J.; Uhen, M.D. 2001. New discoveries of vertebrates from a near-shore marine fauna from the Early Miocene of northwestern Venezuela. Paläontologische Zeitschrift 75: 227-232.
- Sánchez-Villagra, M.; Asher, R.J.; Rincón, A.D.; Carlini, A.A.; Meylan, P.; Purdy, R.W. 2004. New faunal reports for the Cerro La Cruz Locality (Lower Miocene, Northwestern Venezuela). *In* Fossils of the Miocene Castillo Formation, Venezuela: contributions on neotropical palaeontology (Sánchez-Villagra, M.R.; Clack, J.A.; editors). Special Papers in Palaeontology 71: 105-116.
- Sánchez-Villagra, M.; Aguilera, O.; Sánchez, R.; Carlini, A.A. 2010. The Fossil Vertebrate Record of Venezuela of the Last 65 Million Years. *In* Urumaco and Venezuelan Paleontology, The Fossil Record of the Northern Neotropics (Sánchez-Villagra, M.; Aguilera, O.; Carlini, A.A.; editors). Indiana Press University: 19-51.
- Shephard, G.E.; Müller, R.D.; Liu, L.; Gurnis, M. 2010. Miocene drainage reversal of the Amazon River driven by plate-mantle interaction. Nature Geoscience 3 (12): 870-875.
- Simpson, G.G. 1943. Una tortuga del Terciario de Venezuela. Revista de Fomento, Venezuela 5 (51-52): 53-64.

- Simpson, G.G. 1947. A Miocene glyptodont from Venezuela. American Museum Novitates 1368: 1-10.
- Stehlin, H.G. 1928. Ein astrapotheriumfund aus Venezuela. Eclogae Geologicae Helvetiae 21: 227-232.
- Strömberg, C.; Dunn, R.; Madden, R.; Kohn, M.; Carlini, A. 2013. Decoupling the spread of grasslands from the evolution of grazer-type herbivores in South America. Nature Communications 4: (1478) 1-8.
- Toro, M.; Steel, J.R. 2002. A high-frequency sequence study: A Miocene deltaic and estuarine succession in the eastern Maracaibo composite foreland basin, western Venezuela. Bulletin of Canadian Petroleum Geology 50 (1): 3-30.
- Urbani, F.; Mendi, D. 2011. Notas sobre la discordancia del margen sur de la cuenca oligo-miocena de Falcón, estados Lara, Falcón y Yaracuy, Venezuela. Boletín de la Academia de Ciencias Físicas, Matemáticas y Naturales 60: 9-25.
- Useche, C.; Cala, P.; Hurtado, H. 1993. Sobre la ecología de *Brycon siebenthalae* y *Mylossoma duriventris* (Piscis, Characidae), en el Río Cafre, Orinoquia. Caldasia 17: 341-352.
- Verde, M. 2002. Icnología de La Formación Camacho (Mioceno Tardío) del Uruguay. Master dissertation on Zoology (Unpublished). Universidad de la República, Facultad de Ciencias: 126 pp. Uruguay.
- Villamil, T. 1999. Campanian-Miocene tectonostratigraphy, depocenter evolution and basin development of Colombia and western Venezuela. Palaeogeography, Palaeoclimatology, Palaeoecology 153: 239-275.
- Vivas, V.; Macsotay, O. 1995. Formación Tememure: Unidad olistostromica Eoceno Tardío-Oligoceno Temprano en el frente meridional de la Napa Piemontina de Venezuela nor-central. Boletín Geológico, Publicacion Especial 11: 95-113.
- Wesselingh, F.P.; Macsotay, O. 2006. *Pachydon hettneri* (Anderson, 1928) as indicator for Caribbean-Amazonian lowland connections during the Early-Middle Miocene. Journal of South American Earth Sciences 21: 49-53.
- Wesselingh, F.P.; Salo, J.A. 2006. Miocene perspective on the evolution of the Amazonian biota. Scripta Geologica 133: 439-458.
- Weston, E.M.; Madden, R.H.; Sánchez-Villagra, M.R. 2004. Early Miocene Astrapotheres (Mammalia) from Northern South America. *In Fossils of the Miocene* Castillo Formation, Venezuela: contributions on neotropical palaeontology (Sánchez-Villagra, M.R.; Clack, J.A.; editors). Special Papers in Palaeontology 71: 81-97.

- Wheeler, C.B. 1960. Estratigrafía del Oligoceno y Mioceno inferior de Falcón occidental y nororiental. *In* Congreso Geológico Venezolano, No. 3, Memorias 1: 407-465. Caracas.
- Wheeler, C.B. 1963. Oligocene and lower Miocene stratigraphy of western and northeastern Falcón Basin, Venezuela. American Association of Petroleum Geologists, Bulletin 47: 35-68.
- Wilson, J.L. 1975. Carbonate Facies in Geologic History. Springer: 471 p. New York.
- Wright, J.D.; Miller, K. 1992. Miocene stable isotope stratigraphy, Site 747, Kerguelen Plateau. *In* Proceedings of the Ocean Drilling Program, Scientific Results (Wise, S.W.; Schlich, R.; editors). College Station, TX (Ocean Drilling Program) 120: 855-866.
- Zachos, J.; Shackleton, N.; Revenaugh, J.; Palike, H.; Flower, B. 2001. Climate response to orbital forcing across the Oligocene-Miocene boundary. Science 292: 274-278.

Manuscript received: September 10, 2013; revised/accepted: June 03, 2014; available online: June 13, 2014.