

## Early Cretaceous riparian vegetation in Patagonia, Argentina

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**Abstract: Early Cretaceous riparian vegetation in Patagonia, Argentina.** An autochthonous plant assemblage found at the Bajo Grande locality, Anfiteatro de Ticó Formation, Baqueró Group, Early Aptian of Santa Cruz province, Argentina, is described. Sedimentological data show an alternation of sandstone and limestone, with recurrent fossil levels associated to a fluvial system. The frequent flood in areas close to the levee had an influence on the vegetation growing along the banks of the channels. A plant assemblage composed of *Ricciopsis grandensis* nov. sp. (bryophyte), *Adiantopteris tripinnata* nov. sp., Schizaeaceae pinnules (ferns), and the gnetalean fructification *Ephedra verticillata* nov. sp. was found in this paleoenvironment. The taphonomical studies suggest that the plant assemblage was autochthonous, and it grew and was buried in an area near a levee. This result is congruent with environmental data provided by living representative of the fossil taxa analyzed.

**Key words:** Taphonomy, Flood plain, Bryophytes, Ferns, Ephedraceae, Baqueró Group, Early Aptian, Argentina.

The Baqueró Group (early Aptian) occurs in the Nesocratón Deseado basin in Santa Cruz province, Argentina. The deposits are continental, mostly lake and fluvial systems. Stratigraphic studies show this Group to cover volcanoclastic rocks of Jurassic age with angular unconformity. Three formations are recognized within the Baqueró Group: Anfiteatro de Ticó at the base; Bajo Tigre in the middle and Punta del Barco at the top (Cladera *et al.* 2002).

The Baqueró Group flora has been described since 1960, including systematic and biostratigraphy studies (Archangelsky, 1963, 1966, 1967; Menéndez, 1965, 1966; Taylor & Archangelsky, 1985; Archangelsky & Taylor, 1986; Archangelsky & Del Fueyo, 1989; Del Fueyo, 1991; Villar de Seoane, 1998, 1999, 2000; among others). However, the taphonomical and paleoenvironmental studies of this flora are few; A. Archangelsky *et al.* (1995) and Cladera & Cúneo (2002) described for the Punta del Barco Formation, a plant community composed mainly of Araucariaceae, Podocarpaceae, Cycadales and Gleicheniaceae buried by airfall volcanic ash.

The purpose of this paper is to analyse the sedimentology, taphonomy and palaeoecology of a plant community composed of bryophytes, ferns and ephedracean-type fructifications found in a flood plain at a different locality. The Bajo Grande site (Figs. 1, 3A) yielded abundant fossils, including fructifications found isolated (Archangelsky

& Villar de Seoane, 2004) or in organic connection with vegetative parts (Archangelsky, 1968).

### MATERIAL AND METHODS

The specimens were collected at the Estancia Bajo Grande locality in Santa Cruz province, Argentina (Figs. 1, 3A) and referred to the Anfiteatro de Ticó Formation (Baqueró Group). Isotopic dating (Corbella, 2001), confirm that the Anfiteatro de Ticó Formation is 118.56±1.4 Ma old (Early Aptian).

The taphonomic work was done on a 3 m<sup>2</sup> area (Figs. 2, 3B), where the plant assemblage occurs. The fossils correspond to liverwort impressions, filicalean fronds (impressions and compressions with cuticle badly preserved), and branch compressions with reproductive structures of ephedracean affinity. The cuticle of the reproductive structures is strongly carbonized and the treatments performed to recover the membrane showed poor results. The rock maceration gave no palynological data either. Observations were made using light (LM) and scanning electron microscopies (SEM). Classification of Tryon & Tryon (1982) was used for ferns and Taylor & Taylor (1993) for bryophytes and ephedracean fructifications. The specimens are stored in the Paleobotanical Collection of the Buenos Aires Natural Sciences Museum «Bernardino Rivadavia» (BA Pb, BA Pb MEB).

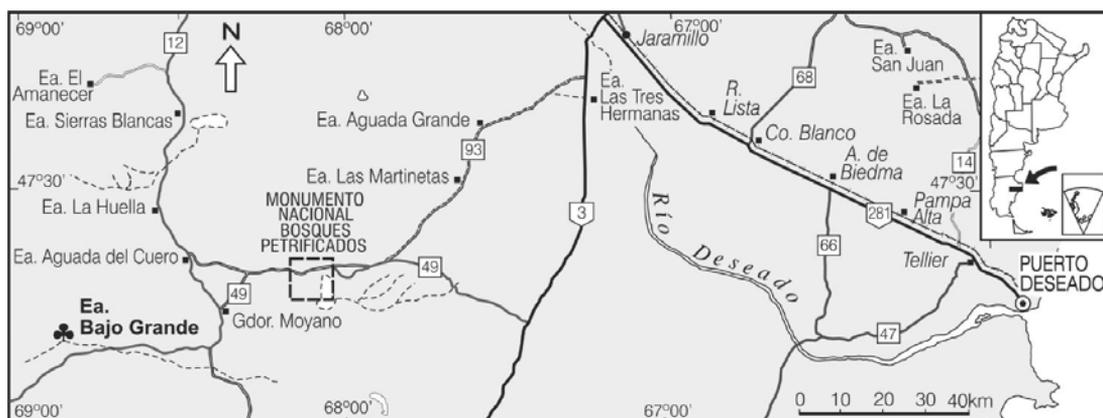


Fig. 1. Location map of the studied area.

## SEDIMENTOLOGICAL AND TAPHONOMICAL ANALYSIS

### Conglomerate and coarse sandstone channel facies

The basal surface of channel deposits is erosive and irregular, forming up to 0,30 m deep steep slopes. The channel basal fill is composed of clast-supported conglomerates with sandy matrix. The most common sedimentary structures are planar and trough-crossed bedding (Fig. 3C) with medium scale heterogeneous sets up to 0,60 m (Fig. 2). Clasts attain a maximum 3 cm size; they are eroded from paleorelief in the underlying Chon-Aike and Bajo Grande formations (Middle Jurassic and Lower Cretaceous). The conglomerates show thickening to coarsening grain arrangements in cycles composed of associated sandstone and rarely limestone up to 1.5 m thick and 40 m wide. These channels move laterally and show a tabular geometry. The thick-medium sized sandstones with isolated clasts show trough-cross bedding structures of a maximum of 0.40 m divided in normal graded sets. These sandstones are usually associated with conglomerates or they form cosets up to 1.5 m thick. Sandstones carry wide branches and trunks 0.20 m in diameter. Fluvial cycles are of Gm-Gt-St type (Miall, 1985). Reactivated fluvial channels show Gm-Gt- or Gt-St cycle types, which would correspond to longitudinal bars, followed by linguoid bars showing transitional contacts on sandstone. Towards the top of the sequence, coarse conglomerate facies disappear and medium sandstones with trough cross bedding dominate. The sandstone thickness is up to 1.5 m and it alternates with siltstone deposits. Paleocurrents indicate a SE flow (135° N).

### Floodplain fine sandstone and siltstone facies

Floodplain deposits are up to 4 m thick and represented by interstratified siltstone and mudstone up to 5 cm thick, massive or laminated, light to dark grey in colour. These beds bear abundant carbonized woody fragments. Fine sandstone beds, up to 10 cm thick, alternate with finer sediments of the floodplain deposits and bear well-preserved plant remains. The internal structure of the sandstones is ripple crossed and laminated with small lenses of mudstone (Fig. 3D). The alternation of sandstone and mudstone beds occurs five times in the floodplain sequence. Paleosols including 1 cm long by 3 mm diameter roots are also commonly found in this alternated sequence. The contact surface between the sandstone beds and finer sediments is both sharp and undulating.

### Interpretation

The conglomerate facies (Gm-Gt) was deposited under low energy conditions (Reineck & Singh, 1980). The St facies can be interpreted as a channel fill (Reineck & Singh, 1980), or else as migration of transversal bars with linguoid crests (Allen 1983; Miall, 1983, 1996). At times this facies tends to fill the deepest part of the channel forming conglomerate-sandstone bars. In some cases, the erosive surfaces are interpreted as a fluctuation in the flow stages. Thus, the channel fill is made of conglomerate-sandstone (Gm-Gt-St), while abandoned facies are represented by siltstone as cycles of Gm-Gt-St-Fm type. Fine-grained cycles are of St-Fm-FI-Fr type, and characterize flood plain deposits laterally associated with channels. This plain was formed by an almost continuous sedimentation of fine sediments by overbank flooding and crevasse

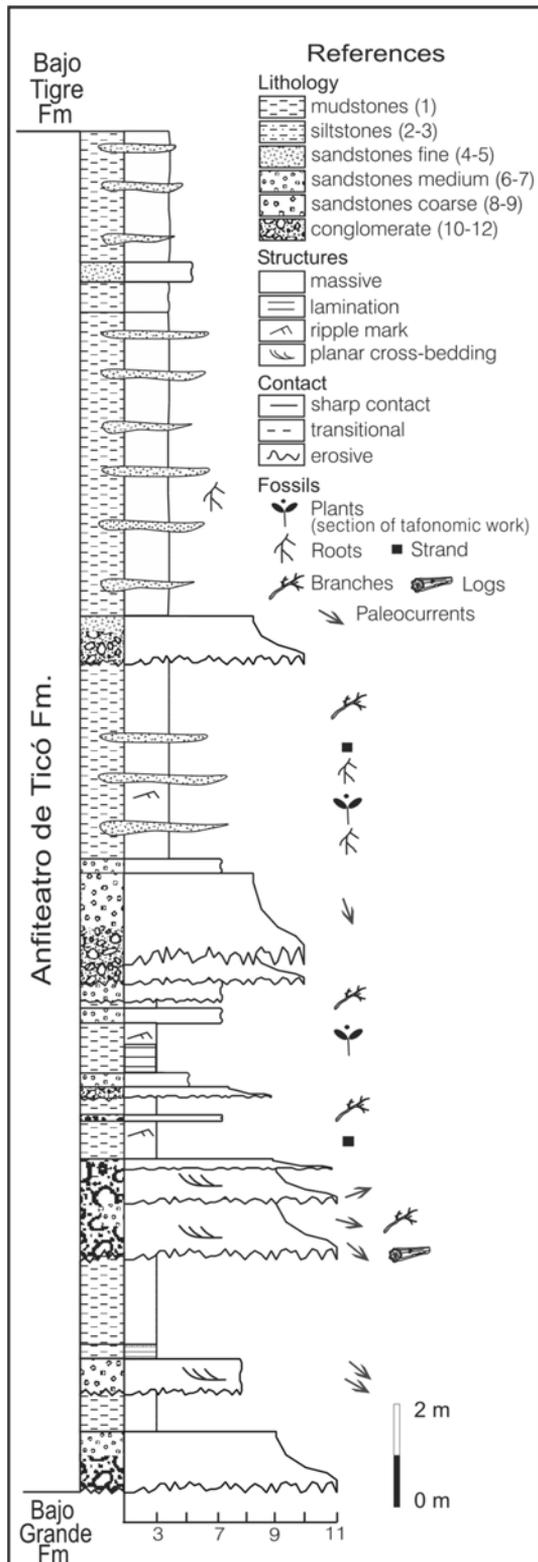


Fig. 2. Stratigraphic section showing location of the fossiliferous beds.

splay represented by sandstone beds. Paleosols indicate interruptions in sedimentation. This facies association may have been part of a levee, as indicated by a high sandstone/ siltstone ratio due to a constant river rising. Based on the characteristics of the associated facies, a low energy of a moderate sinuosity river is suggested.

Plint (1983) has described similar facies for the Middle Eocene of the Hampshire basin in England, while Cladera *et al.* (2002) indicate similar fluvial styles in other areas of the Desierto Mesocretácico.

### Taphonomy

Our observations are based on the 3 m<sup>2</sup> where the plant remains were found. A detailed examination of the surrounding sediments of the fossiliferous area was conducted (both laterally and vertically) but no plant remains were found.

The following taxa were found (percent values show proportion of specimens of each taxon): *Adiantopteris* (49,2%); gnetalean fertile fragments (46,5%); a Schizaeaceae frond (1,54%) and *Ricciopsis* sp. (3,07%).

The fern *Adiantopteris* includes tripinnate fronds without any preferable orientation and crossing the bedding plane (both the sandstones and laminated siltstone), and its pinnule toothed margins and apex are well preserved.

The fragmentary impressions of the ephedracean fertile plant are composed of third order and lower branches with a maximum size of 10 cm long and 0.2 cm wide. Like *Adiantopteris* remains, these fragments are found with no preferred orientation and cross several bedding planes. The margins of the delicate terminal fructifications are perfectly preserved, indicating the absence of water transport. Although these ephedracean fructifications show no evidence of aquatic transport, they are not attached to the vegetative parts.

### SYSTEMATICS

Division HEPATOPHYTA Smith 1972  
Order MARCHANTIALES Smith 1972  
Family RICCIACEAE Smith 1972

Genus *Ricciopsis* Lundblad 1954

Type species. *Ricciopsis florinii* Lundblad 1954

*Ricciopsis grandensis* nov. sp.  
(Fig. 4 A)

*Holotype*. BA Pb 13504.

*Other studied material*. BA Pb 13505.

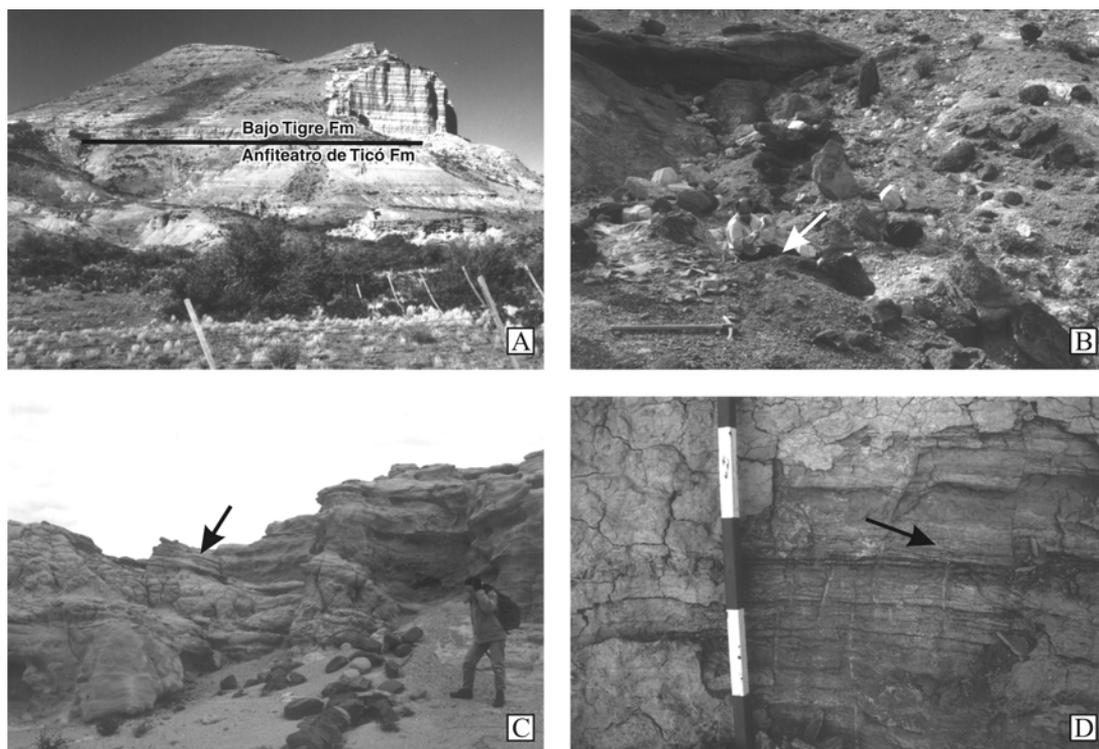


Fig. 3. A-D. Bajo Grande Locality. A. Bajo Grande Locality general view. The line separates beds of the Anfiteatro de Ticó Formation (below) and Bajo Tigre Formation (above). B. Fossiliferous level (white arrow). C. View of a fluvial channel. Arrow shows the through-cross bedding structures. D. Detail of the floodplain where the alternation of sandstone and mudstone beds can be observed. Arrow indicates the ripple cross.

**Locality.** Estancia Bajo Grande, Santa Cruz Province, Argentina.

**Stratigraphic horizon.** Baqueró Group, Anfiteatro de Ticó Formation, Early Cretaceous (Aptian).

**Etymology.** The specific name refers to the fossiliferous locality where the specimens have been recovered.

**Diagnosis.** Plant thalloid. Thalli in groups, forming rosettes up to 1.5 cm in diameter. Thal- lus branched dichotomously up to three times. Branches ranging from 0.10-0.20 cm wide and 0.20 -0.50 cm long. Entire margins, rounded apex and strongly marked midrib.

**Description.** The specimens are represented by impressions belonging to fragments of a ro- sette-shaped thallus up to 1.5 cm in diameter. The thallus has several dichotomous branches which are divided up to three times. It seems that the complete rosette may have had six main branches from the centre. Each branch is linear to wedge-shaped with a strongly marked midrib on the dorsal side, showing entire margins that end in

a rounded apex. The first order branch is 0.15-0.20 cm wide and 0.50 cm long while the second order branch is 0.10-0.20 cm wide and 0.20 cm long and the last order branch is 0.07 cm wide and 0.20-0.40 cm long. Due to preservation no rhizoids were observed.

**Remarks.** The oldest bryophyte megafossil remains in Argentina occur in the Upper Carboniferous of San Juan province, where *Muscites amplexifolius* was described by Ottone & Archangelsky (2001). For the Mesozoic, only one bryophyte was described by Halle in 1913. He studied small fragments of *Marchantites* sp. from mid-Cretaceous sediments of the Lago San Martín (Santa Cruz province). Later Lundblad (1955) assigned these specimens to *M. hallei*.

*Ricciopsis grandensis* represents the first record of Ricciaceae in Argentina. The comparison with other described species from Sweden and Iran (Oostendorp, 1987), shows the following differences. *R. florinii* Lundblad (1954) from the Early Jurassic of Skromberga, Scania province (Sweden), differs by having a larger and crowded rosette-thalli of 2.5 cm in diameter. From

the same locality, Lundblad (*op. cit.*) also recorded *R. scanica* which is smaller (1 cm in diameter) and has four main branches. *R. iranica* Fakhr (1975) from Zhurab locality (Late Triassic-Early Jurassic of Iran), has small star-like thalli (1 cm in diameter) and narrower branches (up to 0.1 cm).

The presence of liverworts in the Baqueró Group is also confirmed through the palynological analyses made by Archangelsky & Villar de Seoane (1996), who studied several spore species of bryophyte, among which two have suggested

*Riccia* affinities (*Naiaditasporea gemmata* Archangelsky & Villar de Seoane and *Coptospora foveolata* Archangelsky & Villar de Seoane).

**Occurrence.** Living hepatics occur in polar, temperate and mainly in tropical areas, where they are found related with shady and moist terrestrial habitats (Smith, 1972). *Riccia* can be terrestrial or aquatic, living on the shore of rivers and brooks. However, certain species are truly aquatics and grow floating on the water surface (Watson, 1971). According to this, the thalli morphology varies from a rosette type to a filiform type, respectively.

Division PTERIDOPHYTA Tryon & Tryon  
1982

Class FILICOPSIDA Tryon & Tryon 1982  
Order POLYPODIALES Tryon & Tryon 1982  
Family PTERIDACEAE Tryon & Tryon 1982

Genus *Adiantopteris* Vassilevskaja 1963 (in  
Tajtadzhan, Vajrameiev & Radchenko 1963)

**Type species.** *Adiantopteris seawardii* (Yabe, 1905) Vassilevskaja 1963 (in Tajtadzhan, Vajrameiev & Radchenko 1963)

***Adiantopteris tripinnata* nov. sp.**  
(Figs. 4 B, C)

**Holotype.** BA Pb 13514 a, b.

**Paratypes.** BA Pb 13509 a, b.

**Other studied material.** BA Pb 13504, BA Pb 13505, BA Pb 13508, BA Pb 13510, BA Pb 13511, BA Pb 13512, BA Pb 13513, BA Pb 13516, BA Pb 13517 a, b, c, BA Pb 13519, BA Pb MEB 186.

**Locality.** Estancia Bajo Grande, Santa Cruz Province, Argentina.

**Stratigraphic horizon.** Baqueró Group, Anfiteatro de Ticó Formation, Early Cretaceous (Aptian).

**Etymology.** The specific name refers to the ramification order of the leaf fragments.

**Diagnosis.** Leaves imparipinnate, bipinnate and tripinnate. Rachises strong and slender. Pin-

nae triangular in shape. Pinnules simple, sub-opposite, small and sessile or sub-petiolate; flabelliform, up to 10 mm long and 2 mm wide. Laminae divided in two to four cuneiform segments with straight margins and dentate apex. Teeth concentration 1 to 2 per millimeter. Only one principal vein radiates and divides dichotomously several times innervating each tooth.

**Description.** The fronds are represented by impressions approximately 6.2 cm long and 2.3 cm wide (Fig. 4 B). The leaves are imparipinnate and bipinnate, but in the specimen BA Pb 13514 a, b, the frond disposition suggests a tripinnate organization (Fig. 4 B). They have strong and slender rachises 4.7 cm long and 0.1 cm wide. The pinnae are up to 2.5 cm long and 1.5 cm wide and have a triangular shape; the pinnules are simple, the basal one being petiolate and the most apical completely sessile (Fig. 4 C). The laminae are divided into two to four segments 5 mm deep and 2 mm wide. The segments are cuneiform, united by only one decurrent base with an insertion angle of approximately 45°. In some specimens, the segments may overlap. They have a dentate apex and straight margin, one is parallel while the other is perpendicular to the rachis. The teeth are separated from each other by a deep U-shape or V-shape valley, ending in an acute apex (Fig. 4 C). The principal vein radiates from the base and divides dichotomously several times, tri to pentafurcate. The concentration is about 3 veins per millimeter. Each vein ends in a corresponding distal tooth.

**Remarks.** *Adiantopteris* was created by Vassilevskaja (in: Tajtadzhan *et al.*, 1963, pp. 586) for Mesozoic-Neogene sterile fronds similar to the living *Adiantum*. Most *Adiantopteris* species were recorded from Jurassic to Miocene beds of Laurasia (Tajtadzhan *et al.*, *op.cit.*), *A. tripinnata* being the first recorded Cretaceous find of Patagonia. The pinnules of *Adiantopteris seawardii* described by Yabe (1905) from the Upper Jurassic-Lower Cretaceous of Korea are larger (15 mm long) and have fewer segments.

*Adiantopteris prigorovskii* Vachrameiev 1952 from the Albian of Kazakhstan has smaller pinnules (7 mm long), the lamina is entire and more expanded in outline. In Patagonia, Berry (1928) illustrated *Adiantum patagonicum* from the Tertiary of Santa Cruz Province. It has isolated and larger (3 cm) stipitate pinnules, and is divided into two terminal and two lateral more or less bisected lobes.

Palynological records from the Anfiteatro de Ticó Formation show the presence of adiantacean spores as *Leiotriletes regularis* (Pflug) Krutzsch (Archangelsky & Villar de Seoane, 1994).

**Occurrence.** *Adiantum* contains about 150 species, widely distributed in pantropical regions. It also extends south to southern South America to Alaska, New Zealand and northeastern Asia, except in areas with extremely cold or dry climates (Tryon & Tryon, 1982).

Family SCHIZAEACEAE Tryon & Tryon 1982

**Genus *Incertae Sedis***  
(Fig. 4 D)

**Studied material.** BA Pb 13514 a and b.

**Locality.** Estancia Bajo Grande, Santa Cruz Province, Argentina.

**Stratigraphic horizon.** Baqueró Group, Anfiteatro de Ticó Formation, Early Cretaceous (Aptian).

**Description.** The material consists of a leaf impression 3 cm long and 1.5 cm wide. The leaf is imparipinnate and has a triangular shape. The rachis is 1 mm wide, slender and straight. The pinnae decrease in size toward the apex, from 9 mm long (the basal) to 2 mm long (the apical) and 3 to 1 mm wide respectively. They are falcate with rounded apex, lobulate margins and asymmetrical base with the adnomic side wider than the catadromic side, that is almost parallel to the rachis. The lobes in the basal pinnae are deeply divided, originating up to four incipient imparipinnate and opposite pinnules. The pinnae have a middle vein and bifurcated secondary veins.

**Remarks.** This leaf impression is included in the Schizaeaceae because its morphology is very similar to the living *Anemia*, especially *A. glareosa* Gardn. that lives in the South American tropics (Tryon & Tryon, 1982, p. 64, fig. 6.12). Among the sterile *Anemia*-type fossils, two genera are known that differ from the specimen here described: *Aneimidium* Schimper, 1869 from the Lower Cretaceous of Germany, that has flabelliform pinnules, and *Aneimites* (Dawson) Ettingshausen, 1865 from the Carboniferous of Germany, with seven to three lobed pinnules.

**Occurrence.** *Anemia* is found in open habitats and well-drained sites of the American tropics, growing on ravine banks, stream borders, shrubby hillsides, small cliffs and sometimes it grows in savannahs, open forests or rain forests (Tryon & Tryon, 1982).

Division GNETOPHYTA Gifford & Foster  
1989  
Order GNETALES Gifford & Foster 1989  
Family EPHEDRACEAE Gifford & Foster  
1989

Genus *Ephedra* Linné 1753

**Type species.** *Ephedra distachya* Linné 1753

***Ephedra verticillata* nov. sp.**  
(Figs. 4 E-I)

**Holotype.** BA Pb 13521 a and b

**Paratype.** BA Pb 13524

**Other specimens studied.** BA Pb13506, 13507, 13515, 13519, 13520, 13522, 13523, 13525 a, b, 13526, 13527, 13528; BA Pb MEB 187, 188 and 189.

**Locality.** Estancia Bajo Grande, Santa Cruz Province, Argentina.

**Stratigraphic horizon.** Baqueró Group, Anfiteatro de Ticó Formation, Early Cretaceous (Aptian).

**Etymology.** The specific epithet refers to the whorl arrangement of branches.

**Diagnosis.** Stems with verticillate branches and seed-bearing structures. Third order branches in whorls of three to six per node. Stems and branches with ridges, up to 0.2 cm wide and 2.2 cm long. Seed-bearing structures single or in clusters of three to five, sessile, rounded to oval in outline and up to 0.15 cm wide and 0.20 cm long. Each structure with two oval-acuminate seeds 0.8 mm wide and 1.8 mm long.

**Description.** The specimens are compressions of fertile fragments represented by straight and strong stems with verticillate branches ending in seed-bearing structures (Fig. 4 E). Most of the material shows up to third order branches occurring in whorls of three to six per node (Fig. 4 F). The stems and branches have longitudinal ridges and no scales or leaves are observed. The stems are 0.07-0.2 cm wide and 1.0-2.2 cm long (incomplete). The first order branches are 0.2 cm wide and vary from 0.5-3.9 cm in length; the second order branches are 0.1-0.2 cm wide and 1.2-2.5 cm long; finally, the paratype is 0.05-0.15 cm wide and 0.4-1.2 cm long. At the end of these branches seed-bearing structures are born, which can be single or in clusters of three to five units (Figs. 4 G, I). These fertile organs are sessile and rounded to oval in outline and up to 0.15 cm wide and 0.20 cm long. Each structure bears two oval-acuminate seeds 0.8 mm wide and 1.8 mm long, which have a striate surface (Fig. 4 H). In some specimens, the seeds are subtended by a pair of bracts up to 2 mm long.

**Remarks.** Mesozoic gnetalean mega-fossil remains occur in few regions. *Ephedra archaeorhynchidosperma* described by Yang *et al.* (2005) from the Barremian Yixian Formation, China, is the

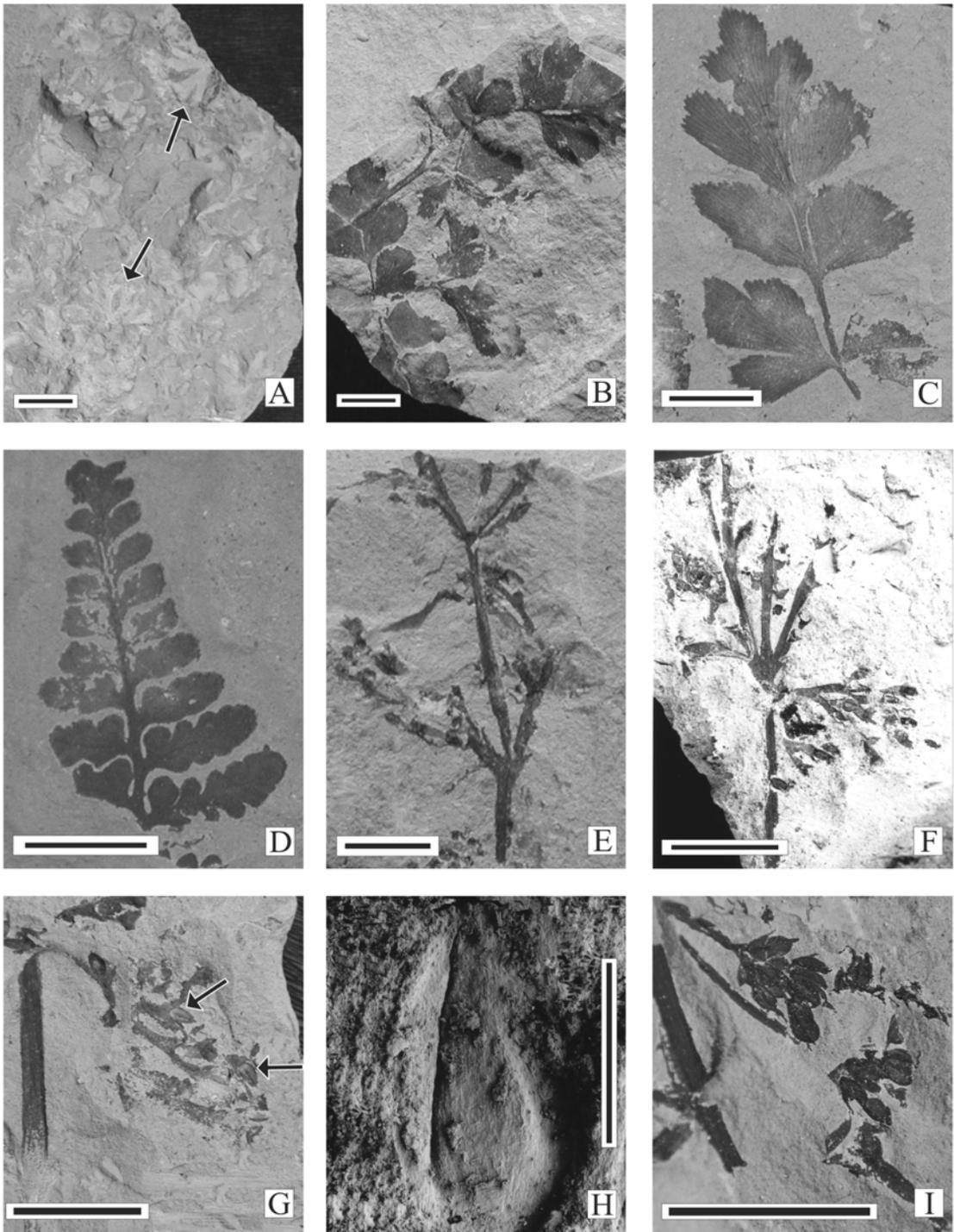


Fig. 4. A- I, Plant assemblage. A. *Ricciopsis grandensis* nov. sp., general aspect of the rosette-shaped thalli (arrows), holotype: BA Pb 13504. B-C. *Adiantopteris tripinnata* nov. sp., B. General view of a frond fragment, holotype: BA Pb 13514 a. C. Detail of a pinna, paratype: BA Pb 13509 b. D. Schizaeaceae, fragment of a pinna, BA Pb 13509 b. E-I. *Ephedra verticillata* nov. gen. et sp. E, F. General aspect of the verticillate branches and seed-bearing structures, holotype: BA Pb 13521 a, BA Pb 13523. G, last order branch with single fertile organs (arrows). H. Detail of a fertile organ. G, H. paratype: BA Pb 13524. I. Clusters of fertile organs, BA Pb 13506.

most similar species to *E. verticillata*. The Chinese specimens differ in having isolated leaves near the nodes and larger seeds (1.5 -4 mm long and 1- 1.6 mm broad) with cuticle preserved. Among other described taxa, the differences with *Ephedra verticillata* are notably marked: *Dinophyton spinosum* Ash (1970) from the Upper Triassic of Arizona has shoots with needle-like leaves (less than 2 mm) with uniovulate lobed cupules 2.6 mm long and 1 mm wide (Krassilov & Ash, 1988). Four different seed-bearing organs from the Middle Jurassic of East Siberia (Krassilov & Bugdaeva, 1988) were characterized by having one ovule ranging from 5- 20 mm long and subtended by axillary bracts. *Drewria potomacensis* from the Early Cretaceous of the Potomac Group of Virginia shows stems with simple and oblong leaves and terminal or axillary reproductive structures containing flattened seeds surrounded by one pair of elliptical ovate bracts (Crane & Upchurch, 1987). A Gnetophyte assemblage from the Early Cretaceous of Transbaikalia presents four new genera of seed-bearing reproductive organs with different morphology that vary from cupules with one-four ovules each to flower-like structures (Krassilov & Bugdaeva, 2000). More recently, coalified seeds closely related with the living *Ephedra* from the Early Cretaceous of Portugal, were described as small seeds between 850 and 1200  $\mu$ m long with a rounded base and pointed micropyle (Rydin *et al.*, 2004).

Palynological data from the Baqueró Group also confirm the presence of the family Ephedraceae through *Ephedripites* (*Equisetosporites*) spp. (Archangelsky & Villar de Seoane, pers. com.).

**Occurrence.** The Ephedraceae live in temperate and tropical regions of Asia, southern Europe and northern Africa. In America, it grows in subtropical regions, in deserts and meadows of the warmer zones (Krüssmann, 1985).

## CONCLUSIONS

The Anfiteatro de Ticó Formation was deposited under fluvial and lacustrine conditions. The sequences of the facies with thickening to coarse grain arrangements and the geometry of tabular bodies, together with lateral displacements, suggest that the fluvial system was meandering.

The fine deposits are associated with accretion deposits due to main flow floods in the plain or as abandoned channel fills. This type of fluvial system is very common in all the areas where the Anfiteatro de Ticó Formation is found (Cladera *et al.*, 2002).

The sandstone and limestone alternation together with the recurrent fossil levels formed by successive flooding of the levee. This was associated with vegetation growing along the banks of the channel, mainly composed of *Adiantopteris tripinnata* and *Ephedra verticillata*, a type of plants that is found only near a levee.

The repetition of this taphocenoses, associated with the levee facies, show a clear cycling scheme of a recurrent vegetation, related to the same sedimentary facies.

The combination of data show that *Adiantopteris tripinnata* and *Ephedra verticillata* were autochthonous, growing and being buried in the area near the levee.

This kind of taphofacies was not found in other sites where the Anfiteatro de Ticó Formation crops out. This unit contains numerous fossiliferous levels that mostly show evidence of transport.

Perhaps, this peculiar topographic setting is responsible of the unique plant assemblage that includes taxa that were not mentioned before, among a large list that includes more than 100 species. In fact the genera *Ephedra*, *Ricciopsis* and *Adiantopteris* are new records for the Anfiteatro de Ticó Formation.

The same section yields abundant fossils in facies in which some transport of plant debris was evident. These allochthonous fossils are very well preserved, however. Fine grained sediments have yielded many fragments of plants, some with their fructifications still attached (*Apterocladus* Archangelsky, 1966; *Androstrobus* Archangelsky & Villar de Seoane, 2004).

The present finding underscores the need for further taphonomic research of this formation and demonstrates that in spite of the large number of taxa that have been described, there is more room for new findings that will improve our knowledge of the Ticó palaeocommunities.

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## BIBLIOGRAPHY

- Allen, J. R. L. 1983. Studies in fluvial sedimentation: bars, bar complexes and sandstone sheets (low - sinuosity braided streams) in the Brownstones (L. Devonian), Welsh Borders. *Sedimentology Geol.* 33: 237 - 293.
- Archangelsky, A., R. Andreis, S. Archangelsky & A. Artabe. 1995. Cuticular characters adapted to volcanic stress in a new cretaceous cycad leaf from Patagonia, Argentina. Considerations on the stratigraphy and depositional history of the Baqueró Formation. *Rev. Palaeobot. Palyn.* 89:213-233.
- Archangelsky, S. 1963. A new Mesozoic flora from Ticó, Santa Cruz Province, Argentina. *Bull. Brit. Mus. (Nat. Hist.), Geol.* 8:45-92.
- 1966. New Gymnosperms from the Ticó Flora, Santa Cruz province, Argentina. *Bull. Brit. Mus. (Nat. Hist.), Geol.* 13: 259- 295.
- 1967. Estudio de la Formación Baqueró, Cretácico Inferior de Santa Cruz, Argentina. *Revista del Museo de La Plata (Nueva Serie) Paleontología*, 5: 63-171.
- 1968. On the genus *Tomaxellia* (Coniferae) from the Lower Cretaceous of Patagonia (Argentina) and its male and female cones. *Bot. J. Linn. Soc.* 61:153-165.
- Archangelsky, S. & G. M. Del Fueyo. 1989. *Squamastrobos* gen. nov., a fertile podocarp from the Early Cretaceous of Patagonia, Argentina. *Rev. Palaeobot. Palynol.* 59: 109-126.
- Archangelsky, S. & T. N. Taylor. 1986. Ultrastructural studies of fossil plant cuticles. II. *Tarphyderma* gen. n., a cretaceous conifer from Argentina. *Amer. J. Bot.* 73: 1577-1587.
- Archangelsky, S. & L. Villar de Seoane. 1994. Estudios palinológicos de la Formación Baqueró (Cretácico), provincia de Santa Cruz, Argentina. VI. *Ameghiniana* 31: 41-53.
- 1996. Estudios palinológicos de la Formación Baqueró (Cretácico), provincia de Santa Cruz, Argentina. VII. *Ameghiniana* 33: 307-315.
- 2004. Cycadean diversity in the Cretaceous of Patagonia, Argentina. Three new *Androstrobus* species from the Baqueró Group. *Rev. Palaeobot. Palynol.* 131: 1-28.
- Ash, S. R. 1970. *Dinophyton*, a problematical new plant genus from the Upper Triassic of south-western United States. *Palaeontology* 13: 646-663.
- Berry, E.W. 1928. Tertiary fossil plants from the Argentine Republic. *Proc.U.S. Nat. Mus.* 73: 1-27.
- Cladera G. & R. Cúneo. 2002. Fossil plants buried by volcanic ash in the Lower Cretaceous of Patagonia. In: De Renzi, M., Pardo Alonso, M. V., Belinchon, M., Peñalver, E., Montoya, P. and Marquez-Aliaga, A. (eds.), *Current Topics on Taphonomy and Fossilization*. Valencia, pp.399- 403
- Cladera, G., R. Andreis, S. Archangelsky & R. Cúneo. 2002. Estratigrafía del Grupo Baqueró, Patagonia (provincia de Santa Cruz, Argentina). *Ameghiniana*, 39:3-20.
- Corbella, H. 2001. Tuffs of the Baqueró Group and the Mid-Cretaceous frame Extra andean Patagonia, Argentina. *11º Congreso Latinoamericano de Geología y 3 Congreso Uruguayo de Geología*. Paper 190 7, 6pp (CD).
- Crane, P. R. & G. R. Upchurch.1987. *Drewria potomacensis* gen et sp. nov., an Early Cretaceous member of Gnetales from the Potomac Group of Virginia. *Am. J. Bot.* 74: 1722-1736.
- Del Fueyo, G. 1991. Una nueva Araucariaceae cretácica de Patagonia, Argentina. *Ameghiniana* 28: 149-161.
- Ettingshausen, C. 1865. *Die Farnkräuter der Jetztwelt* Carl Gerold's Son, Vienna, pp 298.
- Fakhr, M. M. S. 1975. Contribution a l'étude de la flore rhétoliasique de la formation de Shemshak de l'Elbourz (Iran). *Publication du Laboratoire de Paléobotanique de l'Université Paris*. VI, 2 CNRS-A. O.10517.
- Halle, T. G. 1913. Some mesozoic plant-bearing deposits in Patagonia and Tierra del Fuego and their floras. *K. svenska Vetensk Akad. Handl.* 51: 1-58.
- Krassilov, V. A. & S. R. Ash. 1988. On *Dinophyton*-Protognetalean Mesozoic plant. *Palaeontographica B* 208: 33-38.
- Krassilov, V. A. & E. V. Bugdaeva. 1988. Gnetalean plants from the Jurassic of Ust- Balej, East Siberia. *Rev. Palaeobot. Palynol.* 53: 359-374.
- 2000. Gnetophyte assemblage from the Early Cretaceous of Transbaikalia. *Palaeontographica B* 253: 139-151.
- Krüßmann, G. 1985. *Manual of cultivated conifers*. Timber Press. Portland- Oregon. pp. 361.
- Linné, C. 1753. *Species Plantarum*, pp. 1040. Lundblad, B. 1954. Contributions to the geological history of the Hepaticae. Fossil Marchantiales from the Rhaetic-Liasic coalmines of Skromberga (Prov. of Scania), Sweden. *Svensk Botanisk Tidskrift* 48: 383-417.
- 1955. Contributions to the geological history of the Hepaticae. II. On a fossil member of the Marchantiaceae from the Mesozoic plant-bearing deposits near Lago San Martin, Patagonia (Lower Cretaceous). *Botaniska Notiser* 108: 22-39.
- Menéndez, C. A. 1965. *Sueria rectinervis* nov. gen et sp. de la flora fósil de Ticó, Provincia de Santa Cruz. *Ameghiniana* 4: 3-11.
- 1966. Fossil Bennettitales from the Tico Flora, Santa Cruz province, Argentina. *Bull. Brit. Mus. (Nat. Hist.) Geol.* 12:1-42.
- Miall, A. D. 1983. Basin analysis of fluvial sediments. In: Collinson JD, Lewin JL (eds) *Modern and ancient fluvial systems. Spec. Publs. Int. Sediment.* 6: 279-286
- 1985. Architectural element analysis: a new method of facies analysis applied to fluvial deposits. *Earth Sci. Rev.* 22: 261-308.
- 1996. *The geology of fluvial deposits*. Springer-Verlag Berlin-Heidelberg, pp. 582.
- Oostendorp, C. 1987. *The Bryophytes of the Palaeozoic and the Mesozoic*. Bryophytorum Bibliotheca, 34. J. Cramer (ed.). Berlin, pp. 112.
- Ottone, E. G. & S. Archangelsky. 2001. A new bryophyte from the Upper Carboniferous of Argentina. *Ameghiniana* 38: 219-223.
- Plint, A. G. 1983. Facies, environments and sedimentary cycles in the Middle Eocene Bracklesham For-

- mation of the Hampshire Basin: Evidence for global sea-level changes? *Sedimentology* 30: 625-653.
- Reineck, H. E. & I. B. Singh. 1980. *Depositional sedimentary environments, with reference to terrigenous clastics*. 2d, Springer – Verlag, New York, pp. 549.
- Rydin, C., K. R. Pedersen & E. M. Friis. 2004. Early Cretaceous evidence for an ancient origin of *Ephedra*. 7<sup>o</sup> International Organization of Paleobotany Conference. Abstracts, Bariloche, Argentina, pp. 98.
- Schimper, W. P. 1869-74. *Traité de paléontologie végétale ou la flore du monde primitif*, J.B. Bailliere et fils, Paris, 1: 1-740.
- Smith, G. 1972. *Cryptogamic Botany. Bryophytes and Pteridophytes*. Volume II. TMH Edition, New Delhi, pp. 399.
- Tajtadzhan, A. L., V. A. Vajrameiev & G. P. Radchenko. 1963. Algae, Musci, Psilophytes, Lycopodiales, Sphenophytes and Ferns. In: I.A. Orlov (ed.) *Principles of Palaeobotany*, 4. Ac. Cs. URSS. pp. 698. (In russian).
- Taylor, T. N. & S. Archangelsky. 1985. The cretaceous pteridosperms *Rufflorinia* and *Ktalenia* and implications on cupule and carpel evolution. *Amer. J. Bot.* 72:1842-1853.
- Taylor, T. N. & E. L. Taylor. 1993. *The Biology and Evolution of Fossil Plants*. Prentice Hall, New Jersey, pp. 982.
- Tryon, R. M. & A. F. Tryon. 1982. Ferns and Allied Plants. With Special Reference to Tropical America. Springer-Verlag, New York, pp. 857.
- Vachrameev, V. 1952. *Stratigraphy and fossil flora of the Western Kaszastan cretaceous beds*. In: Regional Stratigraphy of URSS, 1, pp. 258 (in russian).
- Villar de Seoane, L. 1998. Comparative study of extant and fossil conifer leaves from the Baqueró Formation (Lower Cretaceous), Santa Cruz Province, Argentina. *Rev. Palaeobot. and Palynol.* 99: 247-263.
- 1999. *Otozamites ornatus* sp. nov., a new benettitalean leaf species from Patagonia, Argentina. *Cret. Res.* 20: 499-506.
- 2000. *Rufflorinia papillosa* sp. nov. from the Lower Cretaceous of Patagonia, Argentina. *Palaeontographica B*, 255: 79-85.
- Watson, E. U. 1971. *The structure and life of Bryophytes*. Hutchinson, London, pp. 211.
- Yabe, H. 1905. Mesozoic plants from Korea. *Journ. Coll. Sci. Univ. Tokyo*, 20: 13-28.
- Yang, Y., B. Geng, D. L. Dilcher, Z. Chen & T.A. Lott. 2005. Morphology and affinities of an Early Cretaceous *Ephedra* (Ephedraceae) from China. *Am. J. Bot.* 92: 231-241.

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