

Albian dinoflagellate cysts from the Kachaike Formation, Austral Basin, Southwest Argentina

M. Verónica GULER¹ & Sergio ARCHANGELSKY²

¹Departamento de Geología, Universidad Nacional del Sur, CONICET. San Juan 670, (8000) Bahía Blanca. E-mail: vguler@criba.edu.ar. ²Museo Argentino de Ciencias Naturales «Bernardino Rivadavia», CONICET. Angel Gallardo 470, C1405DJR Buenos Aires. E-mail:sarcang@fibertel.com.ar

Abstract: The palynological assemblages of the Kachaike Formation at the Bajo Comisión section, Austral Basin, Argentina, are mainly composed by pteridophyte and bryophyte spores, gymnosperm and angiosperm pollen, dinoflagellate cysts and other marine and freshwater algae. This study presents the dinoflagellate cyst assemblages recovered from the lower part of the section. The biostratigraphic index dinoflagellate cysts taxa suggest an early Albian age for the lower part of the Kachaike Formation and the assemblages are attributed to the *Muderongia tetracantha* Interval Zone (Helby). The dominance of well preserved pollen and spores together with the presence of dinoflagellate cysts, suggest that the lower levels of the Bajo Comisión section were deposited in marine shallow waters, in agreement with the sedimentary facies.

Key words: Dinoflagellate cysts, Albian, Austral Basin, Argentina

The Cretaceous deposits in the Austral Basin were grouped in three tecto-sedimentary cycles: Rio Mayer, Lago San Martin and Lago Viedma, ranging from the Berriasian to the Maastrichtian (Arbe, 2002). The Kachaike Formation was included in the Lago San Martin cycle (early Aptian – early Turonian), as the highstand phase within the late Albian – early Turonian Kachaike – Piedra Clavada subcycle. North of Lago San Martín, Cladera & Limarino (in prep.) analyzed exposures of the Kachaike Formation at the Bajo Comisión section (Fig. 1). The 280 m thick integrated sequence consists of sandstones, shales and conglomerates in a vertical arrangement of facies, passing from a marine-deltaic to a moderate to low sinuosity fluvial paleoenvironment (Fig. 2).

The age of the Kachaike Formation is debatable and ranges from late Aptian to early Cenomanian according to different authors. Aguirre Urreta (2002) synthesized the information on lower Cretaceous invertebrate fossil records from Santa Cruz Province. Most of the age-diagnostic molluscan fossils found in the lago San Martin area indicate late Aptian-Albian ages for the formation.

Previous biostratigraphical studies based on megafloora and palynoflora, particularly at the Arroyo Caballo Muerto section (Fig. 1), proposed a late Aptian-early Albian age for the Kachaike Formation (Rebasa, 1982; Gamero, 1982,

unpublished report; Baldoni & Batten, 1991; Baldoni *et al.*, 2001; Cúneo & Gandolfo, 2005). Other palynological analyses in the same section include the systematic study mainly of pollen and spores by Baldoni (1987) and Archangelsky & Llorens (2003; 2005).

In the Bajo Comisión section, Passalía and Archangelsky (2002) recognized six levels bearing fossil remains of diverse gymnosperms and angiosperms, and two angiosperm leaf morphotypes were described by Passalía (2003). Age-diagnostic angiosperm leaves and fossil pollen in the upper part of the Bajo Comisión section suggest a late Albian-early Cenomanian age (Barreda & Archangelsky, 2006).

This study is focused on the lower part of the Kachaike Formation at the Bajo Comisión section where dinoflagellate cysts are present. The biostratigraphical results are mainly compared with those from the Arroyo Caballo Muerto section presented by Baldoni *et al.* (2001).

The aim of this study is to present further evidences of marine components useful for a biostratigraphical analysis, in order to discuss the age of the Kachaike Formation. This is particularly relevant to contribute in the elucidation of palaeobiological events such as the origin and evolution of the angiosperms, since micro and megafloora components of this group in the Kachaike Formation are in constant study.

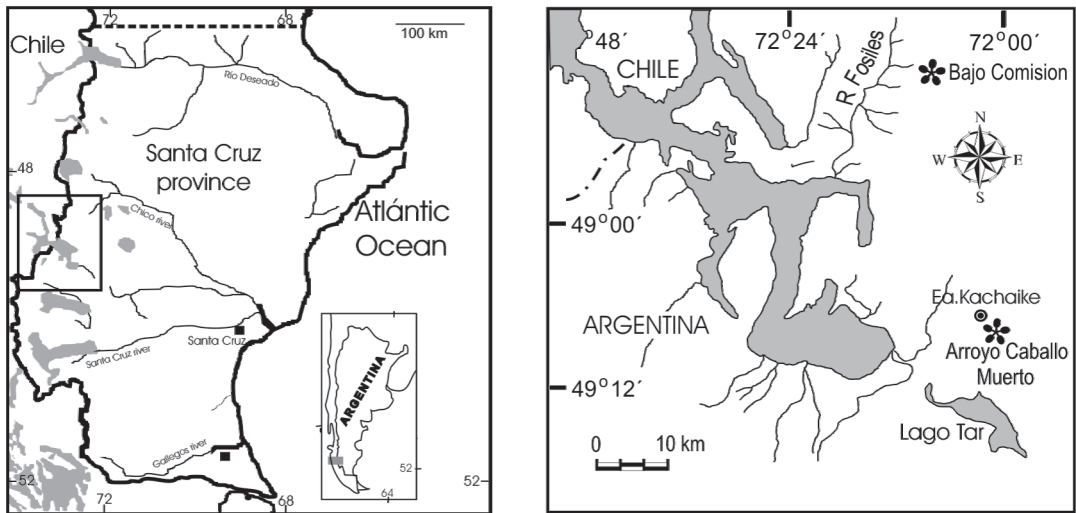


Fig. 1. Map showing location of the Bajo Comisión and Arroyo Caballo Muerto sections in the Lago San Martín area.

MATERIALS AND METHODS

Seventeen samples from the Kachaiké Formation at the Bajo Comisión were processed for palynomorphs. Treatment included hydrofluoric, hydrochloric acid and, for the dinoflagellate cysts, the organic residues were filtered with 25 μm mesh sieves. The residues were stained using Bismarck C and mounted in glycerine jelly.

Light microscopy was undertaken using Nikon Eclipse 600 microscope serial number 772751. Images of specimens were captured by a Nikon Coolpix 950 digital camera. The nomenclature corresponds to Fensome and Williams (2004). Timescale corresponds to Gradstein *et al.* (2004).

PALYNOLOGICAL ASSEMBLAGES AND AGE

The palynoflora from Bajo Comisión is mainly composed by pteridophytes, bryophytes, gymnosperms, angiosperms, dinoflagellate cysts and other remains of marine and freshwater algae. Statistical analysis indicates that the dinocyst / sporomorph ratio is variable throughout the section, and conforms three main intervals that reflect paleoenvironmental changes, from bottom to top: deltaic - marine conditions, subaqueous to subaerial platform and subaerial to fluvial platform (Cladera & Limarino in prep; Archangelsky, personal observation). The transgressive pelites of the lower part of the Bajo Comisión section (PBC10 to PBC15) yielded palynological assemblages with marine elements,

mainly dinoflagellate cysts (Fig. 2). A complete list of taxa determined is given in the Appendix. The dinocyst assemblages are dominated by gonyaulacalean cysts, though peridinialean are well represented. The abundance and diversity of dinocysts are variable, with highest frequencies in the four upper samples (PBC10 to PBC13), reaching up to 30% at PBC10 (Archangelsky, personal observation). The two lower levels bear extremely sparse, poorly-preserved assemblages, almost barren of dinocysts.

The dominance of well preserved pollen and spores together with the presence of dinocysts suggest that the lower deposits of the Bajo Comisión section accumulated in marine shallow waters near the shoreline.

The four lower samples of the Kachaiké Formation at Bajo Comisión bearing dinocysts, indicate an early Albian age. The most important biostratigraphic index species are *Dinopterygium cladoides* (Fig. 3.O,P), *Muderongia tetracantha* (Fig. 3.J) and *Prolixosphaeridium conulum* (Fig. 3.K,L). The presence of *Muderongia tetracantha* indicates an age not younger than early Albian (Helby *et al.*, 1987). The first appearance of *Prolixosphaeridium conulum* is at the mid Albian, according to Morgan (1980) and the base range of *Dinopterygium cladoides* (as *D. tuberculatum*) is considered to be mid Albian (Morgan, 1980; Helby *et al.* 1987). However, *D. tuberculatum* appears to be regularly present in Australia from the base of the Albian (Helby, pers. comm.). The occurrence of *P. conulum* may be a weak stratigraphic marker in these assemblages, since other strong mid Albian markers are

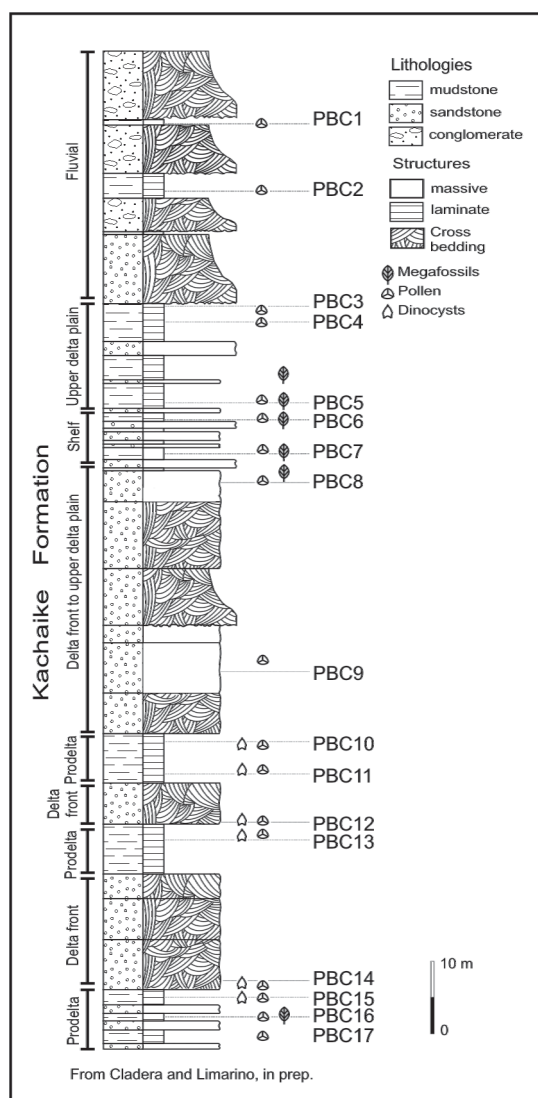


Fig. 2. Integrated stratigraphic section of the Kachaike Formation at the Bajo Comisión.

absent. Therefore, the samples are interpreted to be of early Albian age and attributed to the *Muderongia tetracantha* Interval Zone of Helby *et al.* (1987). However, a mid Albian age for the uppermost levels of the lower part of the section is not totally discarded.

DISCUSSION AND CONCLUSIONS

The Kachaike Formation in the Lago San Martín area appears to extend throughout the Albian based on palynomorphs and other micro and mega fossils recovered at the Bajo Comisión

and Arroyo Caballo Muerto sections. The diagnostic dinoflagellate cysts allow to propose an early age for the lower part of the Kachaike Formation.

In an unpublished report, Gamarro (1982) stated that the Kachaike Formation at the Arroyo Caballo Muerto section was of late Aptian-early Albian age. The author described nine palynologically productive samples and two of them yielded dinoflagellate cysts (98/81 and 689/81); these two dinocyst assemblages were re-studied in Guler and Archangelsky (2003). The lower sample (98/81) bears a scarce and poorly diverse dinocyst assemblage, with long ranging species that impede to state an age for the base of the section. Sample 689/81, from the middle part of the section, contains dinoflagellate cysts with Albian affinities for which a late Albian age was proposed. However, regarding the lack of continuity of the dinocyst records the age of these assemblages must be confirmed in future studies.

Baldoni and Batten (1991) analyzed the megaspores recovered from the Kachaike Formation at the Arroyo Caballo Muerto and proposed a late Aptian-early Albian age. Later, Baldoni *et al.* (2001) stated for the same section, a late Aptian-early Albian (probable early Albian) age, based on the spores, pollen, megaspores and microplankton recovered in 12 of the 14 samples analyzed (2012 to 2015, 2112 to 2119 and a plant level). They stated a minimum early Albian age for the whole section by the presence of the key dinocyst taxon *Muderongia tetracantha* and the absence of angiosperm tricolpate pollen. However, according to the range chart presented in Baldoni *et al.* (2001) *Muderongia tetracantha* is only present in two samples from the lower part of the section. This fact allows to correlate the lower levels of the Kachaike Formation in both Bajo Comisión and Arroyo Caballo Muerto section. Therefore, dinocyst assemblages from the lower part of the Kachaike Formation in both sections would indicate an early Albian age, and suggest at the same time a correlation with the *Muderongia tetracantha* Interval Zone of Helby *et al.* (1987). Otherwise, angiosperm pollen and megafloora recovered from the upper levels of the Bajo Comisión section suggest a younger late Albian age for the upper Kachaike Formation (Barreda & Archangelsky, 2006). The lack of dinocyst records in these continental deposits does not add further support for this dating.

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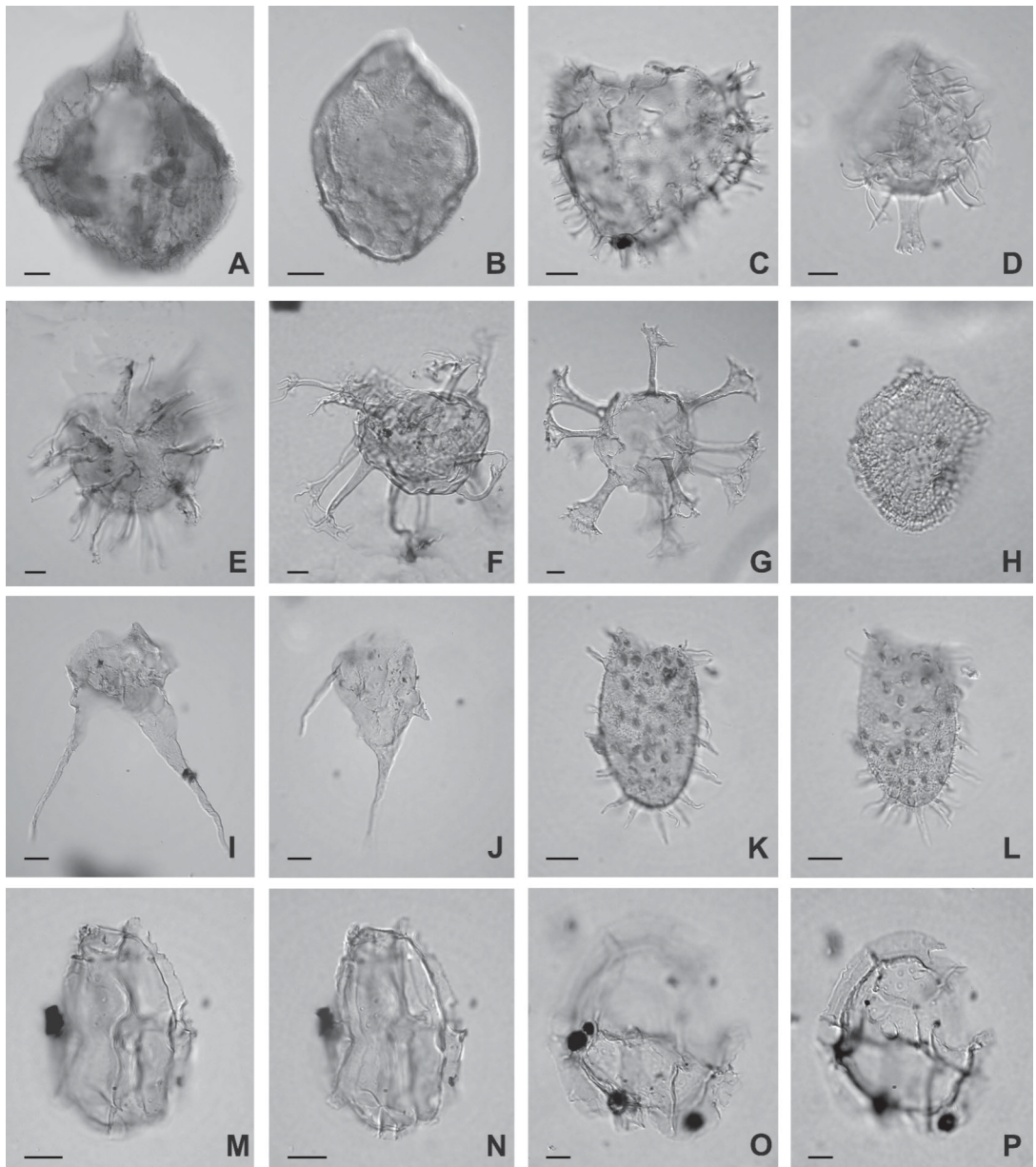


Fig. 3. **A**, *Cribroperidinium orthoceras* (Eisenack) Davey, PBC10E 51,2/103,5; dorsal view, intermediate focus. **B**, *Batiacasphaera* sp. cf. *B. granulosa* Cookson & Eisenack, PBC11C 40/107; dorsal view, low focus. **C**, *Circulodinium distinctum* (Deflandre & Cookson) Jansonius, PBC11C 43/108; dorsal view, low focus. **D**, *Coronifera oceanica* Cookson & Eisenack *emend.* May, PBC11C 36,5/111; right lateral view; high focus. **E**, *Systematophora cretacea* Davey, PBC11E 36,5/113; general view, intermediate focus. **F**, *Oligosphaeridium* sp., PBC12E 25,5/101,1; dorsal view, intermediate focus. **G**, *Oligosphaeridium pulcherrimum* (Deflandre & Cookson) Davey & Williams, PBC10D 46/99,8; oblique apical view, high focus. **H**, *Chlamidophorella nye* Cookson & Eisenack 1958, PBC11C 35/105; general view. **I**, *Odontochitina operculata* (Wetzel) Deflandre & Cookson, PBC10C 27,5/109; dorsal view, high focus. **J**, *Muderongia tetracantha* (Gocht) Alberti *emend.* Monteil, PBC10C 37/109; ventral view, high focus. **K**, **L**, *Prolixosphaeridium conulum* Davey, PBC13C 33/105; dorsal view; **K**, low focus; **L**, high focus. **M**, **N**, *Carpodinium granulatum* Cookson & Eisenack *emend.* Leffingwell & Morgan, 1977, PBC11D 50,5/104,5; dorsal view; **M**, intermediate focus; **N**, low focus. **O**, **P**, *Dinopterygium cladoides* (Eisenack & Cookson) Stover & Evitt, PBC13B 34/104,5; apical-antapical view; **O**, high focus; **P**, low focus.

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Appendix

List of the dinoflagellate cysts encountered in this study. The following species are fully referenced by Fensome & Williams., 2004

- Apteodinium granulatum* Eisenack, 1958 emend. Lucas-Clark, 1987
- Apteodinium* sp.
- Batiacasphaera* sp. cf. *B.granulosa* Cookson & Eisenack, 1974. Fig. 3 B
- Carpodinium granulatum* Cookson & Eisenack 1962b emend. Leffingwell & Morgan, 1977. Figs. 3 M, N
- Chichauoadininium boydii* (Morgan 1975) Bujak & Davies, 1983
- Chlamyphorella «ambigua»* (Deflandre, 1937) Stover & Helby, 1987
- Chlamyphorella nye* Cookson & Eisenack 1958. Fig. 3 H
- Circulodinium distinctum* (Deflandre & Cookson, 1955) Jansonius, 1989 Fig. 3 C
- Coronifera oceanica* Cookson & Eisenack, 1958 emend. May 1980. Fig. 3 D
- Cribroperidinium orthoceras* (Eisenack, 1958) Davey 1969. Fig. 3 A
- Cribroperidinium* sp.
- Dinopterygium cladoides* (Eisenack & Cookson 1960) Stover & Evitt 1978. Figs. 3 O, P

- Florentinia laciniata* Davey & Verdier, 1973
Florentinia mantellii (Davey & Williams 1966)
Davey & Verdier, 1973
Kiokansium unituberculatum (Tasch in Tasch *et al.* 1964) Stover & Evitt, 1978
Muderongia tetracantha (Gocht, 1957) Alberti, 1961 *emend.* Monteil, 1991. Fig. 3 J
Odontochitina operculata (Wetzel, 1933) Deflandre & Cookson, 1955. Fig. 3 I
Odontochitina shinghii Morgan, 1980
Odontochitina sp
Oligosphaeridium complex (White, 1842) Davey & Williams, 1966
Oligosphaeridium pulcherrimum (Deflandre & Cookson, 1955) Davey & Williams, 1966. Fig. 3 G
Oligosphaeridium sp. Fig. 3 F
Prolixosphaeridium conulum Davey 1969. Figs. 3 K, L
Prolixosphaeridium parvispinium (Deflandre, 1937b) Davey *et al.*, 1969
Spiniferites ramosus (Ehremberg, 1838) Mantel, 1954
Stephodium sp.
Systematophora cretacea Davey 1979b. Fig. 3 E
Trichodinium castanea Deflandre, 1935