Radiolarian Age Constraints of Mid-Cretaceous Black Shales in Northern Tunisia

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1. Introduction

Mid-Cretaceous pelagic deposits outcropping in Northern Tunisia include organic-rich beds locally associated with high abundance of radiolarian microfauna, which are interpreted as the signature of the two global oceanic anoxic events OAE1 and OAE2 (Talbi, 1991; Saïdi & Belayouni; 1994; Caron et al., 1999; Amédro et al., 2005, Heldt et al., 2008; Khazri et al., 2009; Soua et al., 2009; Robascynski et al., 2010; Ben Fadhel et al., 2011). Several studies have stated the close association between organic-rich sediments and radiolarian in the Atlantic and Tethyan realms (Marcucci-Passerini et al, 1991; O’Dogherty, 1994; Erbacher & Thurow, 1998; Danelian et al., 2004, 2007).

In North African margins, the radiolarian biostratigraphy have focused upon radiolarian-bearing Jedidi Formation which has been thoroughly discussed by Cordey et al, (2005) and Boughdiri et al, (2007). The first attempts at dating radiolarian series in Northern Tunisia show that radiolarian associated with carbonate-siliceous beds, have yielded useful diagnostic radiolarian assemblages (Cordey et al., 2005; Soua et al., 2006; Ben Fadhel et al., 2010).

Albian and Cenomanian-Turonian black shales of Northern Tunisia were considered to have good generative oil source rock (Layeb, 1990; Saïdi & Belayouni, 1994; Bechtel et al., 1998; Ben Fadhel et al., 2011). In this overall context, the restudy and high-resolution biostratigraphy of Albian black shale beds of Lower Fahdene Formation and C/T cherty beds of organic-rich Bahloul Formation outcropping in Northern Tunisia domain have yielded well-preserved and age-diagnostic radiolarians species. The aim of this paper is to: 1) give new illustrations of radiolarian taxa recovered from albian pelagic deposits of north african margins 2) establish a direct age of black shales using radiolarian assemblages 3) compare the radiolarian assemblages with time equivalent investigated in tethyan and east Pacific domains.

2. Geological setting

The area of investigation is located in Northern Tunisia (Fig. 1). Three sections are selected in this study on the basis of occurrence of organic and radiolarian-rich layers:
1. The Jebel Srassif area (Fig. 1a) is located in the northwestern extremity of the ‘Dome Belt’, a complex structure linked to Triassic extrusions and strike-slip faults. According to Chikhaoui et al., (1991) and Chikhaoui & Turki (1996), the observed structural complexity was the result of the extensive tectonic movement, which led to the extrusion of Triassic evaporites during the Albian–Aptian period. Consequently, halokinetic and tilted blocks movements are responsible for the horst and– graben architecture. The so-called ‘tectonic corners’, described by the previous authors, are induced by the reactivation of strike slip faults during the Tertiary compressive phase. The Jebel Srassif section belongs to the subsiding basin of the Mellegue ‘paleograben’ (Chikhaoui et al., 1991) bordered by two structural highs: Koumine to the west and Nebeur to the east. Cretaceous successions are characterized by a thick pile of Aptian to Campanian pelagic sequences, which are affected by multiple non-depositional unconformities and condensed layers (Chikhaoui, 1988).

2. The Fadeloun-Garci-Mdeker structure (fig.1b) in which belongs the Jebel Garci section, is composed of three anticlines, trending North South and considered as the northern prolongation of the N-S axis (Saadi, 1990). The anticlines are separated from the Atlasic domain by the Zaghouan thrust, which its North-eastern part becomes south-verging, commonly defined as the Chérichira-Kondar thrust (Khomsi et al., 2004). The Cretaceous sedimentation was under the control of syn-sedimentary faults trending N140-160 reflected by chaotic and gravitational deposits (Saadi, 1991). Early Cretaceous successions show northward, reduced thickness and affected by hiatus and extreme condensations in Hammam Zriba (Saadi, 1990). The motion of a corridor trending north-south by N140-160 faults has led to the compartmentalization of the seafloor in losangic basins (Saadi, 1991). During the Valanginian – Barremian time span, theses basins were supplied by siliciclastic deposits while condensed sedimentation occupied uplifted horsts (Biely et al., 1973; Saadi et al., 1994).

3. The Oued Kharroub section (Fig. 1c) is located in the Atlas domain (Northern and Central Tunisia), characterized by various facies of Cenomanian-Turonian transition (C-T) deposits, including benthic fauna-rich carbonates e.g. Zebag Formation by Burollet (1956) and Gattar Formation by Boltenhagen and Mahjoub (1974) and organic-rich black shales with pelagic fauna (e.g. Bahloul Formation by Burollet, 1956) respectively of shelf and slope in the southern margin of the Tethyan realm. During this period and since the Jurassic, this domain has been influenced by the opening of the Tethyan palaeosea, its deepening as well as its southern margin migration. Generally speaking, the Bargou area, connected palaeogeographically to central Tunisia, is characterized by (1) emerged palaeohighs displaying gaps and discontinuities (Turki, 1985) and (2) subsiding zones affected by deep-water sedimentation. This area is dominated by N140° and N70° trend faults limiting several blocks. Cretaceous sedimentation varies on both sides. Its structural evolution may be summarized as follow : (1) during the late Jurassic to early Cretaceous, the area was subjected to a major extensional phase that delimited horst and graben systems (Martinez & Truillet, 1987) (2) In the uppermost Aptian, a regional compressional pulsation affecting the north-African platform had resulted from a transpressional scheme (Ben Ayed & Viguier, 1981) (3) New NNE-SSW trend anticline structures appeared attested by the Albian Fahdene Formation onlap features on the reeal aptian Serj deposits in subsurface (Messaoudi & Hammouda, 1994) or upper
Aptian - Albian unconformity in outcrops (Ouahchi et al., 1998). (4) During the Albian, the geodynamic evolution is marked by the sealing of lower Cretaceous structures during an extensional phase that persisted to form graben systems promoting organic-rich and siliceous strata deposition throughout upper Cenomanian to Lower Turonian times (Soua et al., 2009). The major faults in this area are represented by N140° and N70° trend features. The Bahloul thickness is significantly variable in this area. It may varies from 10m to 40m in thickness (Layeb & Belayouni, 1989; Soua & Tribovillard, 2007). Uniquely, in this area, the top of the Bahloul represents many cenomanian olistolith levels (Soua et al., 2006) marking syndepositional tectonic activities (Turki, 1985).

Fig. 1. Geological map of the studied sections (After Chikhaoui et al., 1991; Meddeb, 1986)

3. Result

3.1 Jebel Garci section
The condensed section of Jebel Garci (Fig. 2) begins with orbitolinids-rich green to gray clay alternating with discontinuous sandy limestone beds which are attributed to the Hameima Formation. The clay intervals have also provided fragments of rudist and bryozoans (GA1). The upper part contains olistoliths deposits that gradually pass to a reefal limestone which is outlined at the top by burrowed hardground.

The next successions (GA9 - GA23) which correspond to the “Allam” Member consist of centimeter-thick grey to dark laminated limestone bed and organic-rich black marl intervals.
Upwards, the succession becomes rhythmic and the marly intervals increase in thickness in opposition to limestone beds. The microfauna content yields depauperate planktic foraminiferal assemblages and radiolarian rich microfauna. Fractures related to a strike slip fault outlining the black hales unit are onlapped by a marly intervals and gray limestone beds alternation (GA24-GA27).

The organic-rich beds (GA17 – GA 23) have released a moderately to well-preserved and age-diagnostic radiolarian species. Twenty nine species were recorded in the studied section. The radiolarians appear with few discrete taxa within GA6 level. It provides an assemblage composed of Holocryptocanium barbui Dumitrica, Spongostichomitra elatica (Aliev), Pseudeucyrtis hanni (Tan), Archodictyomitra vulgaris Pessagno, Thanarla brouweri (Tan), Stichomitra simplex (Smirnova and Aliev), Angulobracchia portmanni Baumgartner, Thanarla pacifica Nakaseko and Nishimura. They become diversified and abundant within GA7. It yields an association of Dictyomitra aff. gracilis (Squinabol), Dictyomitra communis Squinabol, Dictyomitra montisserei (Squinabol), Pseudodictyomitra lodogaensis Pessagno, Thanarla praeveneta Pessagno, Archaeodictyomitra aff. A. vulgaris Pessagno, Hiscocapsa sp., Thanarla aff.pulchra (Squinabol), Spongostichomitra elatica (Aliev), Thanarla brouweri (Tan), Angulobracchia portmanni Baumgartner, Stichomitra simplex (Smirnova and Aliev), Stichomitra communis Squinabol.

GA15 sample provided very diversified and abundant radiolarian population. It is composed by Dictyomitra montisserei (Squinabol), Holocryptocanium barbui Dumitrica, Pseudodictyomitra lodogaensis Pessagno, Stichomitra simplex (Smirnova and Aliev), Pseudeucyrtis hanni (Tan), Diacanhocapsa sp., Hiscocapsa grutterinki (Tan) Angulobracchia portmanni Baumgartner, Stichomitra communis Squinabol, Pseudodictyomitra paronai (Aliev), Cryptamphorella conara (Foreman).

GA18 provided Dictyomitra gracilis (Squinabol), Thanarla conica (Squinabol)

The upper part of black shales (G17-23), composed by rhythmic bundles of limestone and marl beds, is characterized by a decrease of radiolarian abundance. The sample GA20 has released a radiolarian assemblages composed of Thanarla brouweri (Tan), Spongostichomitra phalanga O’ Dogherty, Pseudodictyomitra paronai (Aliev), Dictyomitra communis Squinabol, Holocryptocanium barbui Dumitrica, Pseudodictyomitra lodogaensis Pessagno, Dictyomitra gracilis (Squinabol), Dictyomitra montisserei (Squinabol), Spongostichomitra elatica (Aliev), Pseudodictyomitra paronai (Aliev).

Although the uppermost beds have yielded (GA24-27) benthic foraminiferal-rich assemblages, we identified well-preserved radiolarian population (GA24) composed of Pessagnobrachia rara (Squinabol), Stichomitra communis Squinabol, Dictyomitra montisserei (Squinabol), Cryptamphorella conara (Foreman), Holocryptocanium barbui Dumitrica, Pseudodictyomitra lodogaensis Pessagno, Torculum coronatum (Squinabol), Xitus spicarius (Aliev), Obeliscoites vinassai (Squinabol), Hiscocapsa asseni (Tan), Thanarla pulchra (Squinabol), Dictyomitra communis Squinabol, Hiscocapsa grutterinki (Tan).

Marly interval of the top GA27 have released an assemblage of Holocryptocanium barbui Dumitrica, Stichomitra simplex (Smirnova and Aliev), Pseudodictyomitra paronai (Aliev), Dactyliosphaera maxima (Pessagno).

3.2 Jebel Srassif section

The base of Jebel Srassif section (Fig. 3) which constitutes the “Marnes Moyennes” Member, consists of 130 meter-thick alternations of grey marl and limestone, which become dark and laminated at the top. A cyclic marl/limestone bundles (10m) can be distinguished having an
Fig. 2. Jebel Garci section
organic and radiolarian-rich mudstone texture. It is capped by a thick organic-rich limestone bed (20 m) characterized by bituminous odor and yellowish color in patina. This level corresponds to the Mouelha member (Burollet, 1956). The 40 meters of the top consist of an alternation of grey limestones and dark grey-ochre marls yielding septarian nodules characterizing the Defla member. They are overlain by a succession of lenticular limestone beds and grey marl of Azreg member (50 m).

Three samples were selected, based on the good preservation of the faunal assemblages. Among 35 radiolarian morphotypes, only 23 species were figured. Biostratigraphic analysis of the fossil record and planktic foraminifer’s zones (Fig. 3, and Plates 1 and 2) correlation allow us to distinguish the following three radiolarian assemblages:

1. The sample 37 has provided a diversified radiolarian fauna with the co-occurrence of Dictyomitra montisserei (Squinabol), Obeliscoites perspicuus (Squinabol), Tubulustrium transmontanum O’Dogherty, Dictyomitra gracilis (Squinabol), Holocryptocanium barbui Dumitrica, Stichomitra aff. navalis O’Dogherty, Cryptamphorella conara (Foreman), Torculum dengoi (Schmidt-Effing), Stichomitra communis Squinabol, Torculum coronatum (Squinabol), Distylocapsa micropora (Squinabol), Patellula verteroensis (Pessagno), Godia concava (Li & Wu).

2. Radiolarian assemblage recovered from sample 62 is highly diversified at the top of Mouelha blackshales. Likewise, it records an acme of species belonging to Hagiastridae and Cavaspongiidae taxa. This interval shows the co-occurrence of Dispongotripus acutispinus Squinabol, Dactyliosphaera maxima (Pessagno), Pessagnobrachia sp., Cavaspongia euganea (Squinabol), Cryptamphorella conara (Foreman), Pessagnobrachia rara (Squinabol), Dorypyyle communis (Squinabol), Pseudodictyomitra paronai (Aliev), Pseudodictymitra sp., Torculum coronatum (Squinabol), Holocryptocanium tuberculatum Dumitrica, Distylocapsa micropora (Squinabol), Obeliscoites perspicuus (Squinabol), Dactyliosphaera acutispina Squinabol, Dictyomitra gracilis (Squinabol), Thanarla spoletoensis O’Dogherty, Dactyliosphaera lepta (Foreman), Patellula verteroensis (Pessagno), Savaryella novadensis (Squinabol), Savaryella quadra (Foreman), Pessagnobrachia fabianii (Squinabol), Stichomitra communis Squinabol, Holocryptocanium barbui Dumitrica, Xitus aff. spicularius (Aliev), Torculum coronatum (Squinabol), Crolanium aff. spineum (Pessagno),

3. Sample 68 is characterized by the abundance of cryptocephalic nassellaria (Holocryptocanium). Moreover, we notice the first occurrence and bloom of Mallanites triquetrus. This interval shows the co-occurrence of Xitus mclaughlini (Pessagno), Hexapyramis pantanelli Squinabol, Mallanites triquetrus (Squinabol), Thanarla spoletoensis O’Dogherty, Dictyomitra montisserei (Squinabol), Godia concava (Li & Wu), Cryptamphorella conara (Foreman), Torculum coronatum (Squinabol), Cavaspongia euganea (Squinabol), Distylocapsa micropora (Squinabol), Dactyliosphaera maxima (Pessagno), Holocryptocanium barbui Dumitrica, Dactylidiscus longispinus (Squinabol), Dispongotripus acutispinus Squinabol.

3.3 Oued Kharroub section

The outcrop (Fig. 4) is composed mainly by dark clayey limestone and organic-rich black shales with abundant planktic foraminifera. These organic-rich deposits include siliceous beds with abundant radiolarians, an equivalent to “Livello Bonarelli” bed marker (Marcucci Passerini et al, 1991; Salvini and Marcucci Passerini, 1998; Premoli-Silva et al, 1999; Scopelliti et al, 2004; Musavu-Moussavou et al, 2007)
Fig. 3. Jebel Srassif section
Plate 1. 1 – *Dictyomitra gracilis* (SQUINABOL), scale bar: 50m, sample 62. 2 – *Dictyomitra montisserei* (SQUINABOL), scale bar: 50m, sample 68. 3 – *Tubilustrium transmontanum* O’DOGHERTY, scale bar: 50m, sample 37. 4 – *Holocryptocanium barbui* DUMITRICA, scale bar: 50m, sample 37. 5 – *Stichomitria aff. navalis* O’DOGHERTY, scale bar: 50m, sample 37. 6 – *Cryptamphorella conara* (FOREMAN), scale bar: 50m, sample 37. 7 – *Mallanites triquetrus* (SQUINABOL), scale bar: 100m, sample 68. 8 – *Dictyomitra gracilis* (SQUINABOL), scale bar: 100m, sample 62. 9 – *Xitus aff. spicularius* (ALIEV), scale bar: 100m, sample 62. 10 – *Dispongrotius acutispinus* SQUINABOL, scale bar: 100m, sample 62. 11 – *Holocryptocanium tuberculatum* DUMITRICA, scale bar: 50m, sample 62. 12 – *Savaryella quadra* DUMITRICA, scale bar: 100m, sample 62. 13 – *Patellula verteroensis* (PESSAGNO), scale bar: 150m, sample 37. 14 – *Dactyliosphaera lepta* (FOREMAN), scale bar: 50m, sample 62. 15 – *Dactyliodiscus longispinus* (SQUINABOL), scale bar: 50m, sample 68. 16 – *Torculum dengoi* (SCHMIDT-EFFING) Scale bar:
50m, sample: 37. 17 – *Pessagnobrachia* sp., scale bar: 100m, sample 62. 18 – *Cavaspongia euganea* (SQUINABOL), scale bar: 100m, sample 62. 19 – *Stichomitra communis* SQUINABOL, scale bar: 50m, sample 62. 20 – *Dactyliosphaera maxima* (PESSAGNO), scale bar: 100m, sample 62. 21 – *Godia concava* (LI & WU), scale bar: 100m, sample 37. 22 – *Torculum coronatum* (SQUINABOL), scale bar: 150m, sample 62. 23 – *Croatium aff. spineum* (PESSAGNO), scale bar: 100m, sample 62. 24 – *Obeliscoites perspicuus* (SQUINABOL), scale bar: 100m, sample 62.

Plate 2. 1 - *Dictyomitra gracilis* (SQUINABOL), 100µm, GA18. 2 - *Pseudoeucyrtis hanni* (TAN), 100µm, GA15. 3 - *Dictyomitra montisserei* (SQUINABOL), 100µm, GA21. 4 - *Pseudodictyomitra lodogaensis* PESSAGNO, 100µm, GA21. 5 - *Thanarla praeveteta* PESSAGNO, 100µm, GA8. 6 - *Archeodictyomitra aff. vulgaris* PESSAGNO, 100µm, GA8. 7 - *Hisocapsa sp.* 100µm, GA8. 8 - *Dictyomitra communis* (SQUINABOL), 100µm, GA8. 9 - *Thanarla aff. pulchra,* (SQUINABOL), 100µm, GA8. 10 - *Holocryptocanium barbui* DUMITRICA, 100µm, GA20. 11 - *Dictyomitra gracilis* (SQUINABOL), 200µm, GA9. 12 - *Thanarla brouweri* (TAN), 100µm, GA20.
A total of twenty five of radiolarian species are recognized belonging to nassellarians and spumellarians with maximum of eighteen (18) species in sample OKS 11. Their differential stratigraphical range and relative abundance allow to distinct two successive assemblages (R_I and R_II) through the C-T transition. Although, the studied radiolarian species do not exhibit a good potential for biostratigraphic dating, the section is calibrated either by foraminifers and ammonites.

The R_II assemblage spans the upper part of the OAE-2 interval and the organic-poor deposits overlying this interval. It is characterized by a decrease trend of the nassellarian relative abundances (from 87% to 42%). Therefore, maybe dissolution or bad preservation conducted to the absence of this group close to the base of the upper half of the section, across the OKS40-OKS45 samples interval. Many species show rapid and gradual disappearing following a stepwise-like pattern (e.g. Guttacapsa sp., Spongistichomitra elatica, Novixitus sp., Stichomitra stocki, Mita gracilis, Pseudodictyomitra pseudomacrocephala, Thanarla pacifica D. montisserei).

About the associated spumellarians, several species from the R_I assemblage persisted more or less long time (e.g. Archaeocenosphaera aff. vitalis, Crucella messinae, Praecenocaryomma lipmanae, Rhopalosyringium hispidum Pyramispongia glascockensis Pessagno., Cavaspongia euganea (Squinabol), C. Californiensis Campbell and Nishimura, Pseudoeucyrtis spinosa (Squinabol), Archaeocenosphaera ? mellifera O’Dogherty,). Nevertheless, very few species of nassellarians first occurred across the upper half part of the studied section. All these species are represented by dwarf and poorly preserved specimens.
Plate 3. 1 - *Thanarla conica* (SQUINABOL), 100µm, GA18. 2 - *Diacanthocapsa* sp. 200µm, GA15. 3 - *Cryptamphorella conara* (FOREMAN), 50µm, GA22. 4 - 8 - *Pseudodictyomitra paronai* (ALIEV), 100µm, GA20. 5 - *Torculum coronatum* (SQUINABOL), 200µm, GA21. 6 - *Stichomitri simplex* (SMIRNOVA et ALIEV), 100µm, GA15. 7 - *Angulobracchia portmanni* BAUMGARTNER, 100µm, GA15. 8 - *Hiscocapsa* aff. *grutterinki* (TAN) 50µm, GA23. 9 - *Pessagnobrachia rara* (SQUINABOL), 100µm, GA27. 10 - *Xitus spicularius* (ALIEV), 100µm, GA25. 11 - *Spongostichomitra elatica* (ALIEV), 50µm, GA20. 12 - *Stichomitria communis* SQUINABOL, 50µm, GA8.

4. Discussion

Detailed analysis of radiolarian assemblages allows us to attribute a biostratigraphic framework for the organic-rich beds. In the following section, we used zonal scheme proposed by O’Dogherty (1994) for tethyan realms. The age-diagnostic assemblages are discussed and compared with time equivalent investigated in adjacent tethyan domains.
O’Dogherty (1994) proposed a radiolarian zonation for the Albian based on Unitary Associations. He described for the Upper Albian to the base of the Cenomanian the Spoletensis zone divided into three radiolarian subzones: the Romanus, Missilis and Anisa subzones. Bak (1995) established a radiolarian zonation (H. barbui – H. geysersensis) for the Albian–Cenomanian of northern Tethyan domains, based on the co-occurrence of Holocryptocanium barbui DUMITRICA, Holocryptocanium geysersensis PESSAGNO, Novixitus weyli SCHMIDT-EFFING, Squinabollum fossile (SQUINABOL), Cryptamphorella macropora DUMITRICA, Hemicryptocapsa tuberosa DUMITRICA.

The first appearance of B. breggiensis, recorded within Upper Albian basal intervals, coincides with first appearance of radiolarian species Tubilustrium transmontanum O’DOGHERTY, which is confined with the upper part of the Romanus subzone (O’Dogherty, 1994). An assemblage containing Stichomitra navalis and Torculum coronatum was recorded also within this subzone.

Babazadeh & de Wever (2004) described a radiolarian assemblage yielding the co-occurrence of Dictyomitra gracilis, Holocryptocanium barbui and Dictyomitra montisserei and assigned it to Middle–Late Albian age. Nevertheless, the presence of T. dengoi, whose first appearance coincides with the Missilis – Anisa subzones boundary (O’Dogherty, 1994), allows rejuvenating the assemblage age.

Samples recovered from the succession overlying the Mouelha Member blackshales show an assemblage composed of Cryptamphorella conara Dumitrica, Pessagnobrachia sp., and Thanarla spoletensis O’DOGHERTY, which correspond to the lower part of the Appenninica zone and the middle part of the Anisa subzone of O’Dogherty (1994). Although the coexistence of D. lepta, Stichomitra communis and Patellula verteroensis is assigned to early Late Cenomanian age (Erbacher, 1998), this assemblage possibly characterize the Late Albian taking into account the presence of D. maxima whose last occurrence is coeval with the base of Anisa subzone (O’Dogherty, 1994).

Samples recovered from basal beds (GA2-GA6) show high abundance of Pseudodictyomitra lodogaensis and contain some early Cretaceous taxa from Turbocapsula Zone such as A.portmanni and Th. pacifica (O’ Dogherty, 1994; Erbacher and Thurow, 1998; Danelian et al., 2007; Michalik et al., 2008). Thus, a late Aptian age of these beds could not hitherto be ruled out.

According to Erbacher & Thurow (1998), the first occurrence of Pseudodictyomitra lodogaensis coincides with the upper part of G. algerianus Zone. Its last occurrence coincides with the Aptian-Albian boundary and the first occurrence of Mita gracilis (= Dictyomitra gracilis). This taxon is also reported from the Albian to Cenomanian deposits of the Atlantic domain, California and Pacific realms (Thurow, 1988; Karminia, 2006; Palechek et al., 2010).

It is possible that black shale unit of Jebel Garci could underlines the Aptian-Albian boundary. In fact, Danelian (2008) have reported the presence of Thanarla praeveneta from the Upper Aptian – Lower Albian bed which occurs in GA7 beds underlying the black shale successions.

On the other side, Slazcka et al., (2009) described an assemblage containing Angulobracchia portmanni Baumgartner, Dictyomitra communis (Squinabol), Hiscocapsa asseni (Tan), Pseudodictyomitra lodogaensis Pessagno, Pseudoeucyrtis hanni (Tan), almost similar to GA7 taxa. These authors attributed the assemblage to Costata zone that is confined to UA6-9 biochronozones of mid to late Aptian age (O’Dogherty, 1994).

It is noteworthy to point the coexistence of Albian species in all samples such as D.montisserei and D.gracilis with Aptian taxa particularly in GA7, GA 15 and GA26.
In that score, an assemblage recovered from Mid Cretaceous outcrops of Northern Tethys margins was described by Danelian et al., (2007), shows the co-occurrence of P. lodogaensis, Dictyomitra gracilis, Thanarla brouweri, Archaeodictyomitra aff. vulgaris assigning it to the early Albian UA10-11 biochronozone. Danelian et al (2004) consider that an early Albian age of Dercourt Member cannot be ruled out despite the presence of Angulobracchia portmanni and pseudoeucyrtis hanni characteristic of U.A.9. These species are observed hitherto within assemblage from GA15, associated with Dictyomitra montisserei.

Kurilov & Vishnevskaya (2011) described an assemblage extracted from Early Cretaceous outcrops of Pacific domain that does not differ from GA21. It contains Thanarla brouweri, Pseudodictyomitra paronai, Pseudodictyomitra lodogaensis, Holocryptocanium barbui, Dictyomitra cf. montisserei, Dictyomitra communis, and Dictyomitra gracilis indicating an early Albian age.

The sample GA26 has provided an assemblage characterized by high abundance of Hiscocapsa asseni, co-occurring with D.gracilis and D.montisserei. It lies with the UA10 biochronozone of Romanus zone (O’Dogherty, 1994; Danelian et al., 2004).

We suggest that lower part of black shale intervals could be assigned to the upper part of Costata zone (GA5 - GA14) based on the presence of Aptian taxa (i.e. Angulobracchia portmanni, Pseudoeucyrtis hanni). The lower part of this zone coincide with the first occurrence of Microhedbergella praeplanispira planktic foraminifera. Whereas the top coincide with the last occurrence of Angulobracchia portmanni and Pseudoeucyrtis hanni associated with a relative increase in abundance of Archaeodictyomitreae and Williriedellidae families.

The Romanus zone (GA14 - GA27) show the dominance of high diversified nassellarian species. The assemblage recovered from GA17 is composed of Thanarla brouweri, Archaeodictyomitra montisserei, Thanarla conica which is attributed to the middle Albian Mallanites romanus subzone (U.A. 10 -11 biochronozone) (O’Dogherty, 1994; Danelian et al, 2004). However, the first occurrence of Ticinella primula planktic foraminifera is recorded 24 m above GA17 bed. Thus, we suggest that lower part of Romanus zone may be attributed to the Early Albian.

Studies on Cenomanian - Turonian boundary interval show that deposition of radiolarian, organic-rich sediment and large positive carbon isotopic excursion are coeval with extreme fertility conditions and correspond to a large-scale proxy that indicate a hypersiliceous period (Premoli Silva et al, 1999; Racki & Cordey, 2000). The Bonarelli equivalent in Tunisia is commonly known by the Bahloul Formation (Burollet, 1956). In the Bargou area, the Bahloul Formation shows organic-rich intervals interbedding cherty and radiolarian limestone layers (Layeb and Belayouni 1999, Soua and Tribovillard, 2007).

Although the C/T boundary interval outcropping in the Tunisian realm was extensively studied by planktic foraminifera and ammonite biostratigraphy (Maamouri et al, 1994; Nederbragt and Fiorentino, 1999; Abdallah et al., 2000; Amédro et al, 2005), radiolaria assemblages have provided a useful tool for age calibration and subdivision of C/T organic-rich beds in this study.

Two black shale levels were identified in Oued Kharroub section:
1. The first lies with the lower part of Withinella archaeocretacea planktic foraminifera zone, above the highest occurrence of Rotalipora cushmani.
2. The second coincides with the middle part of Heterohelix moremani zone.

The calibration of these levels is based on age-diagnostic radiolarian recovered from biosiliceous limestone beds (Fig. 4).
The OSK 24 yields an assemblage composed of *Rhopalosyringium radiosum* O’Dogherty, *Praecoccaryonna lipmanae* Pessagno, *Acaeniotyle vitalis* O’Dogherty *Rhopalosyringium hispidum* O’Dogherty. The three first taxa have been described by Bak et al (2005) and attributed them to the late Cenomanian – early Turonian. Erbacher (1998) attribute *Rhopalosyringium radiosum* to the early Turonian, but later Musavu-Moussavou and Danelian (2006) expand its range to late Cenomanian. The assemblage contains *Xitus picenus* Salvini and *Marcucci - Passerini* which its range do not exceed the Silviae Zone of Bonarelli (O’Dogherty, 1994; Salvini & Marcucci-Passerini, 1998). Consequently, we assign the lower black shale beds (OKS 11) to the late Cenomanian and to upper part of *Silviae Zone* [U.A 18 biochronozone of O’Dogherty (1994)].

Many authors have stated the occurrence of *Archaecenosphaera mellifera* O’Dogherty within Turonian strata of Boreal and northern Tethyan domains (Bandini et al., 2006; Smreckova, 2011). In East Pacific domain, this taxon, associated with *C. californiensis* and *Pyramidispongia glascockensis* PESSAGNO, is recorded within the Silviae Zone of late Cenomanian age (Bragina, 2009). Salvini & Marcucci-Passerini (1998) stated that *C. californiensis* occurs only in the base of upper assemblage C of Bonarelli Level which lies with the base Superbum Zone defined by O’Dogherty (1994). In the Atlantic domain, the last occurrence of *C. californiensis* is recorded in the late Cenomanian just beneath the organic-rich beds related to the OAE2 (Musavu-Moussavou and Danelian, 2006). Taking into account the paleogeographic similarities between northern and southern Mediterranean Tethys margins, the radiolarian assemblage recovered from OKS44 level could be correlated with upper assemblage (Superbum Zone) of Bonarelli level in Central Italy. Thus, the second black shale lie with the upper part of *Biacuta subzone* of late Cenomanian age, if we take into consideration the position of turonian *Watinoceras* spp. ammonite (Amédro et al., 2005)

5. Conclusion

Biostratigraphic investigations of Albian and C/T boundary intervals in Northern Tunisia show that organic-rich beds are generally associated with high abundance of radiolarian fauna.

Age constraint of organic-rich sediments is established and correlated with biochronozones of O’Dogherty (1994). In the light of these results, we deduce that:

1. Black shale interval of Jebel Garci which is embedded within the “Allam” Member is assigned to the early Albian U.A.10 biochronozone. However, the latest Aptian could not be excluded for the lower part.
2. Late Albian organic-rich beds of Jebel Srassif including cyclic limestone/marl beds of “Marnes Moyennes” and Mouelha Members lie with the boundary interval between U.A. 13 and U.A. 14 biochronozones.
3. Two black shale levels embedded within Bahloul Formation are probably of late Cenomanian age and confined with the U.A.18 biochronozone. The first occurrence of turonian *Watinoceras* spp. ammonite is recorded 70 cm above the second black shale bed (OSK40)

It seems that distribution of radiolarian assemblages of albian and cenomanian-turonian boundary intervals shows some difference from those of Atlantic and east Pacific domains. Preservation index and range discrepancies of some radiolarian species could affect the subdivision resolution. Further studies on radiolarian distribution assemblages and relationships with environmental changes during Mid-Cretaceous time are needed to establish paleogeographic reconstructions of southern tethyan margins.
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