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## Uppermost Albian biostratigraphy and chronostratigraphy

Robert W. SCOTT <sup>1</sup>

**Abstract:** The Albian Stage is the highest chronostratigraphic unit of the Lower Cretaceous Series and underlies the Cenomanian Stage of the Upper Cretaceous Series. The Albian is divided into three substages, each of which is composed of two or three zones based on distinctive and phylogenetically related ammonite assemblages. The uppermost zone of the Upper Albian Substage, the *Stoliczkaia dispar* Zone, is found in many Western European condensed sections. The ammonite assemblage in the thin glauconitic sandstone near La Vraconne, Switzerland, was defined as the 'Vraconnian Stage' in 1868. However this concept has been little used and was abandoned in 1963 as part of the Cretaceous chronostratigraphic scale. A recent proposal to resurrect and redefine this stage is based on a number of criteria and very detailed and reliable stratigraphic data. A quantitative biostratigraphic analysis of the ammonite ranges in the key sections shows that the proposed subzones of the *S. dispar* Zone have discordant ranges. Furthermore, the utility of a 'Vraconnian Stage' between the Albian and Cenomanian stages is geographically limited and the concept embraces one of many depositional sequence cycles of the Albian. The reinstatement of a 'Vraconnian Stage' is not recommended.

**Key Words:** Albian; Vraconnian; ammonites; planktic foraminifers; graphic correlation; age calibration.

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**Résumé :** *Biostratigraphie et chronostratigraphie de l'Albien sommital/terminal.*- L'étage Albien constitue l'unité chronostratigraphique la plus élevée du Crétacé inférieur et repose sous l'étage Cénomaniens du Crétacé supérieur. L'Albien est divisé en trois sous-étages, chacun comprenant deux ou trois zones établies sur des associations d'ammonites distinctes mais phylogénétiquement reliées. La zone sommitale du sous-étage Albien supérieur, la Zone à *Stoliczkaia dispar*, a été identifiée au sein de nombreuses séries condensées en Europe occidentale. L'association d'ammonite reconnue dans le mince niveau de grès glauconieux des environs de La Vraconne (Suisse) a été définie comme 'étage Vraconnien' en 1868. Toutefois ce concept a été peu usité et fut abandonné en 1963 en tant qu'unité de l'échelle chronostratigraphique du Crétacé. Il a été récemment proposé de réhabiliter et de redéfinir cet étage sur la base d'un certain nombre de critères et de données stratigraphiques très détaillées et fiables. Or une analyse biostratigraphique quantitative des répartitions des ammonites dans les coupes clefs révèle que les sous-zones proposées pour la subdivision de la Zone à *S. dispar* correspondent à des intervalles non concordants. En outre l'intérêt de placer un 'étage Vraconnien' entre les étages Albien et Cénomaniens apparaît géographiquement limité, et le concept correspond à une seule des nombreuses séquences de dépôt de l'Albien. Il n'est donc pas recommandé de restaurer un 'étage Vraconnien'.

**Mots-Clefs :** Albien ; Vraconnien ; ammonites ; foraminifères planctoniques ; corrélation graphique ; calibration des âges.

### Introduction

Recently AMÉDRO (2002, 2008), AMÉDRO & ROBASZYNSKI (2008) and ROBASZYNSKI *et alii* (2007) proposed to reinstate the 'Vraconnian Stage' between the Albian and Cenomanian stages. This proposition, which would significantly modify the Cretaceous geologic column, merits careful and thorough analysis. Our objectives are to provide a testable global biostratigraphic database with which to evaluate the ranges of key uppermost Albian

species and to calibrate their ranges to a numerical time scale.

The Cretaceous System is composed of twelve stages that can be correlated world-wide and are of different durations (HANCOCK, 2003). The Albian Stage is the youngest chronostratigraphic unit of the Lower Cretaceous Series and one of the longest Cretaceous stages, about 13 to 15 myr. The Albian Stage was defined by d'ORBIGNY (1840-1842) to include the fossil assemblages in shale and

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glauconite sands cropping out along the Aube River in the Department of Aube on the eastern margin of the Paris Basin (MAGNIEZ-JANNIN & RAT, 1980). A stratotype section along the Aube River was composited from outcrop sections and nearby boreholes (LARCHER *et alii*, 1965; AMÉDRO, 1992). The Albian biota in the Paris Basin is transitional between Boreal and Tethyan realms. The concepts of Albian ammonite zones have evolved over a period of time beginning in 1868 when the "*Ammonites mammillaris*" zone (de RANCE, 1868) (now the

*Douvilleiceras mammillatum* Zone) and the "zone of *Ammonites inflatus*" (de LAPPARENT, 1868) (now considered equivalent to the *Stoliczkaia dispar* Zone) were proposed (RAWSON *et alii*, 1978). The current zonal scheme (Fig. 1) was composed by SPATH (1923) and BREISTROFFER (1947) and modified by OWEN (1971) and AMÉDRO (1992). The zonal succession has been relatively stable since 1947 and most are interval zones defined by the first appearance of an ammonite species (FO; AMÉDRO, 1992).

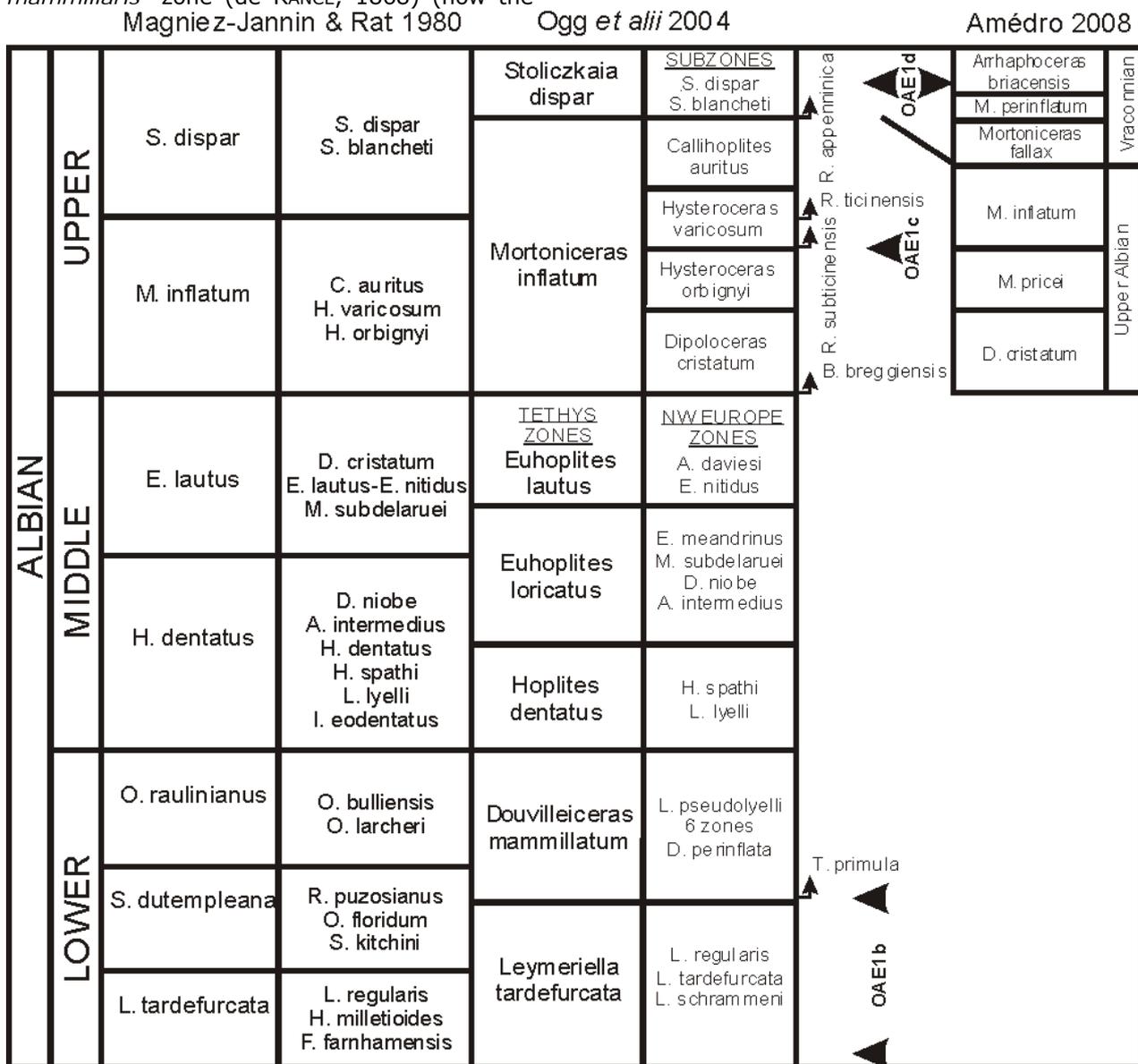


Figure 1: Chart comparing three stage and substage concepts of the Albian (P. DESTOMBES, 1979 in MAGNIEZ-JANNIN & RAT, 1980; OGG *et alii*, 2004, and AMÉDRO, 2008). The positions of Albian anoxic events and key microfossils were interpolated by OGG *et alii* (2004).

The Albian Stage is divided into three substages, Lower, Middle and Upper. Although stratotype sections for these substages are yet to be agreed upon, here they are used formally following HART *et alii* (1996) and are capitalized. The zonal boundaries of the substages have been used consistently since 1947 (Fig. 1). Prior to 1947, however, two different criteria were used to define the base of the Upper Albian

Substage. SPATH (1923, 1941) placed the boundary at the top of the *Dipoloceras cristatum* Subzone, which directly overlies the *Euhoplites lautus* Zone (OWEN, 1971). Alternatively BREISTROFFER (1947) placed the Cristatum Subzone in the basal Upper Albian *Mortonicerases inflatum* Zone. This later opinion has been followed since (BIRKELUND *et alii*, 1984; OWEN, 1984a, 1984b; HANCOCK, 1991; HART *et alii*,

1996; RAWSON & HOEDEMAEKER, 1999; HOEDEMAEKER & RAWSON, 2000; HOEDEMAEKER *et alii*, 2003). The rationale for this change is reviewed by OWEN (1971) and HART *et alii* (1996). In North America a regional transgressive unconformity coincides with the top boundary of the Cristatum Subzone, which is a widespread mappable contact (SCOTT *et alii*, 2003).

The Albian Stage comprises the evolutionary origins and/or diversification of seven families of ammonites. Zones of the Lower, Middle and Upper Albian substages are based mainly on species of the families of Lyelliceratidae, Hoplitidae, and Brancoceratidae. Four heteromorph families first appear in the Albian, Anisoceratidae, Baculitidae, Hamitidae, and Turrilitidae, but are not used to define zones until their appearance in the Upper Cretaceous.

The 'Vraconnian Stage' was proposed by RENEVIER (1868) for a 2 m-thick condensed interval of green glauconitic sand with a distinctive ammonite fauna between the Upper Albian and Lower Cenomanian substages near La Vraconne in western Switzerland. The ammonite assemblage is part of the *Stoliczkaia dispar* Zone and the lithostratigraphic unit correlates with the Upper Gault and Upper Greensand in England (Fig. 1). This stage was discarded in 1963 (COLLIGNON, 1965; RAWSON *et alii*, 1978; HANCOCK, 1991, 2003, among many others). However the 'Vraconnian Stage' has been resurrected and redefined by AMÉDRO (2002, 2008), AMÉDRO & ROBASZYNSKI (2008) and by ROBASZYNSKI *et alii* (2007). AMÉDRO cites five reasons for recognizing 'Vraconnian' sedimentary strata as a stage between the Albian and Cenomanian stages: (1) the interval is mappable in Western Europe, (2) the interval is a third-order depositional cycle that records an important eustatic event, (3) the interval has a distinctive and diverse fossil assemblage that can be recognized outside Europe, (4) its 2 to 3 myr duration is equivalent to that of the Santonian Stage, and (5) in the Vocontian Basin the interval is more than 100 m thick, which is thicker than the underlying part of the Albian (AMÉDRO, 2002, 2008). AMÉDRO has presented detailed lithostratigraphic and biostratigraphic data of twelve sections in Europe, Tunisia, Madagascar, and California to support his proposal. For the first time he presents a regional lithostratigraphic correlation of many of these sections. HANCOCK (2003) reviewed the history of the rejection of the 'Vraconnian' concept by the community of Lower Cretaceous stratigraphers beginning in 1963 and believed that reasons to revive 'Vraconnian' were "trivial". He noted that most 'Vraconnian' sections are condensed intervals.

As a framework for discussing the wisdom of revising the Albian Stage by reinstating the 'Vraconnian Stage', a review of the stage concept is relevant. "A stage is a chronostratigraphic unit of smaller scope and rank than a series. It is most commonly of greatest use in intra-continental classification and correlation, although it has the potential for worldwide recognition" (NACSN, 2005, p. 1582). As a chronostratigraphic unit the Albian Stage has synchronous boundaries and is the physical evidence or 'material referent' of a time interval, the Albian Age. The Albian Stage has traditionally been defined by a set of ammonite biozones (Fig. 1): d'ORBIGNY, 1840-1842; SPATH, 1923; P. & J.-P. DESTOMBES, 1965; OWEN, 1984a, 1984b; HANCOCK, 1991; HART *et alii*, 1996, among others. The Global Stratotype Section and Point (GSSP) have yet to be selected to define the basal boundary although an excellent section has been proposed (KENNEDY *et alii*, 2002). The upper boundary of the Albian Stage is defined by the base Cenomanian Stage GSSP at Mont Risou, France (GALE *et alii*, 1996; KENNEDY *et alii*, 2004).

### Methodology of integrating biostratigraphic data

The integrated ranges of select ammonites in seven key sections defined three zones of the Vraconnian (AMÉDRO, 2008, Fig. 4 ; Fig. 1, Table 1). The ranges of key planktic foraminifera in the Tunisian sections and the French Mont Risou section were added to that set of species by ROBASZYNSKI *et alii* (2007). However most of the ammonite species are found in only one or two sections (AMÉDRO, 2008). To test the accuracy of the integration of species ranges the quantitative technique of graphic correlation is used in this report. Graphic correlation (GC) provides an objective method to compare the ranges of species in multiple sections and the outcome can be tested independently.

Graphic correlation (GC) is a quantitative, non-statistical, technique that determines the coeval relationships between two sections by comparing the ranges of event records in both sections (CARNEY & PIERCE, 1995). A graph of any pair of sections is an X/Y plot of the FOs (first appearances) and LOs (last appearances) of taxa found in both sections. The interpreter places a line of correlation (LOC) through the tops and bases that are at their maximum range in both sections. This LOC is the most constrained hypothesis of synchronicity between the two sections and extends the fewest bioevents. The LOC also accounts for hiatuses or faults at stratal discontinuities indicated by the lithostratigraphic record. The position of the LOC is defined by the equation for a regression line. Explanation and examples of the graphic technique are illustrated by MILLER (1977) and CARNEY & PIERCE (1995). By iteratively graphing

successive sections a database of species ranges is compiled. The accuracy of these ranges depends on the number of sections, preservation and correct identification of the species. Such a database is testable and the

process is transparent so that the fossil occurrence in each section can be evaluated to determine its accuracy. This process compiles data of many specialists who have studied numerous global sections.

| Mont Risou, France, Section    | FO meters | LO meters | Harchies, Belgium                 |         |         |
|--------------------------------|-----------|-----------|-----------------------------------|---------|---------|
| Base limestone interval        | 330 m     |           | Anisoceras perarmatum             | -87.1   |         |
| Actinoceras sulcatus           | 50        | 78        | Anisoceras pseudoelegans          | -95.5   |         |
| Actinoceras subsulcatus        | 25        |           | Callihoplites vracensis           | -87.1   |         |
| Anisoceras salei               | 78        |           | Hamites virgulatus                | -98.4   |         |
| Anisoceras perarmatum          | 78        |           | Hyphoplites subfalcatus (ID cf.)  | -86.1   |         |
| Arrhaphoceras briacensis       | 305       |           | Lepthoplites cantabrigiensis      | -102.1  |         |
| Dipoloceras cristatum          | 18        | 25        | Pleurohoplites subvarians         | -87.1   |         |
| Hysteroeras orbigny (as sp.)   | 78        |           | <b>Strépy, Belgium, Outcrop</b>   |         |         |
| Lechites moreti                | 155       |           | Callihoplites seeleyi             | 30      |         |
| Mantelliceras mantelli         |           | 205       | Callihoplites pulcher             | 30      |         |
| Mariella gresslyi              | 155       |           | Callihoplites tetragonus          | 20      |         |
| Mortoniceras perinflatum       |           | 205       | Cantabrigites subsimplex          | 30      |         |
| Rota globotruncanoides         | 295       |           | Hyphoplites valbonnensis          | 20      |         |
| Turrillitoides hugardianus     | 155       |           | Lechites gaudini                  |         | 20      |
| <b>Folkestone, UK, Section</b> |           |           | Mortoniceras fallax               | 20      |         |
| Base cenomanian ammonites      | 38        |           | Neophlyticeras blancheti          | 20      |         |
| Inoc concentricus              | 0         | 10.5      | <b>ANDRA MAR 203 Core, France</b> |         |         |
| Actinoceras sulcatus           | 9.7       | 13        | Hamites virgulatus (ID cf.)       | -523.23 |         |
| Anisoceras perarmatum          | 26        | 38        | Hyphoplites coelonotus            | -776.75 | -523.38 |
| Arrhaphoceras substuderi       | 26        | 38        | Hyphoplites falcatus              | -455.85 |         |
| Callihoplites auritus          | 14.5      | 24.5      | Hyphoplites valbonnensis          | -487.17 |         |
| Callihoplites cantabrigense    | 25        |           | Lechites gaudini                  | -487.17 |         |
| Callihoplites leptus           | 25        |           | Mariella bergeri (ID cf.)         | -467.25 |         |
| Callihoplites tetragonus       | 26        | 38        | Ostlingoceras puzosianum          | -561.54 |         |
| Callihoplites vracensis        | 26        | 38        | Pleurohoplites renauxianus        | -581.8  |         |
| Hyphoplites coelonotus         | 25        |           | Schloenbachia varians             | -430.8  |         |
| Lepthoplites falcoides         | 26        | 38        | <b>Diégo core, Madagascar</b>     |         |         |
| Mortoniceras inflatum          | 14.5      | 25        | Lechites gaudini                  | -50     |         |
| Pleurohoplites renauxianus     | 26        | 38        | Mantelliceras mantelli            | -23     | -5      |
| <b>Merstham, UK</b>            |           |           | Mariella bergeri                  | -20     |         |
| Anisoceras picteti             | 4.7       | 5         | Neostlingoceras carcitanense      | -23     | -5      |
| Callihoplites seeleyi          | 7         | 7.2       | Scaphites simplex                 | -54     |         |
| Callihoplites vracensis        | 4.7       | 7.2       | Sciponoceras roto                 | -23     | -5      |
| Idohamites elegantulus         | 4.7       | 5         | Stoliczkaia dispar                | -50     |         |
| Lechites gaudini               |           | 7         | Biti breggiensis                  | -215    | -118    |
| Lepthoplites falcoides         | 4.7       | 5         | Planomalina buxtorfi              | -106    | -10     |
| Lepthoplites pseudoplanus      | 4.7       | 7.2       | Planomalina praebuxtorfi          | -118    | -90     |
| Mortoniceras alstonensis       | 4.7       | 5         | Rota appenninica                  | -131    | -10     |
| Mortoniceras fallax            | 4.7       | 5         | Rota brotzeni                     | -23     | -10     |
| Neophlyticeras blancheti       | 4.7       | 6.6       | Rota globotruncanoides            | -23     | -10     |
| Ostlingoceras puzosianum       | 7         | 7.2       | Tici praeticinensis               | -215    | -180    |
| Pleurohoplites renauxianus     | 7         | 7.2       | Tici primula                      | -236    | -135    |
|                                |           |           | Tici subticinensis                | -215    | -155    |
|                                |           |           | Tici ticinensis                   | -180    | -106    |

**Table 1:** Biostratigraphic species and ranges in meters of each section from AMÉDRO (2008).

## Integrated Upper Albian chronostratigraphy

### Process of Graphic Correlation Experiment

To begin the GC experiment the Mont Risou section, southeastern France (GALE *et alii*, 1996; Fig. 2) was selected as the standard reference section because it records continuous basin deposition at a uniform rate of accumulation and it yields diverse ammonites, inoceramids, planktic foraminifera, and nannofossils. This section was cross plotted to itself setting its thickness in meters as the relative time scale. Subsequently seven other sections, which yielded precise and abundant biostratigraphic data, were plotted to it through multiple rounds and the ranges of the fossils extended to the thickness scale at Mont Risou. The first section

plotted was the Kalaat Senan, Tunisia (AMÉDRO, 2008, Fig. 20), which spans from uppermost Albian to Cenomanian; this section also represents uniform deposition. The result is that the scale was extended from the base of the Upper Albian to the Cenomanian/Turonian contact.

The second graphed section was the Diégo well, N Madagascar (AMÉDRO, 2008, Fig. 22). This 300 m cored interval is mainly marl that spans an interval from lowermost Cenomanian to middle Albian (Fig. 3D). The section yields several key planktic foraminifera and in three samples a few ammonites. The LOC of the graph is constrained by a small number of first occurrences and it extends the bases of many foraminifera. The section appears to record continuous Late Albian deposition.

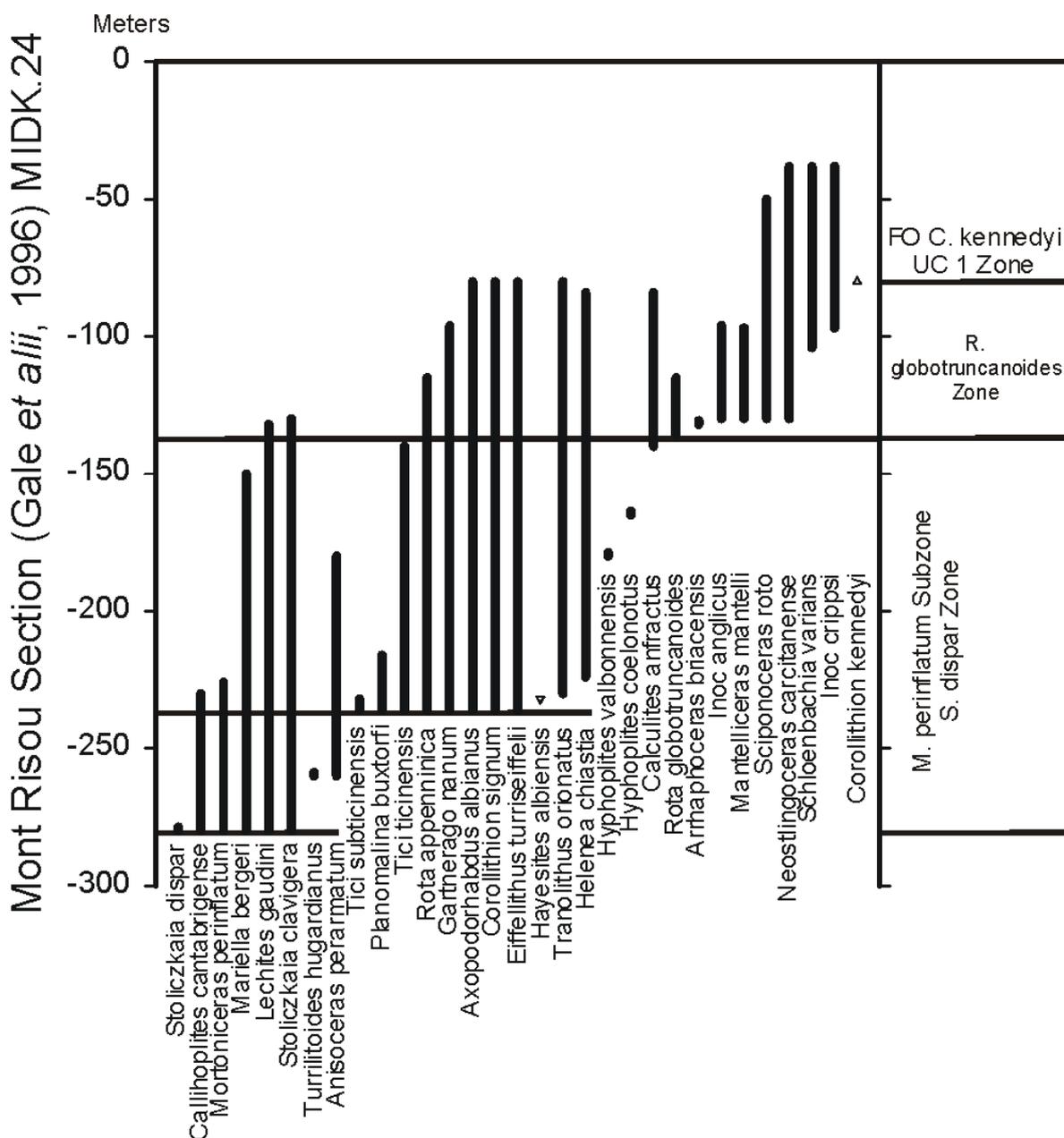


Figure 2: Biostratigraphic range chart of Mont Risou Albian-Cenomanian reference section showing key ammonites, planktic foraminifera and nannofossils (from GALE *et alii*, 1996).



The next section plotted was the Gault Clay in the Folkestone section in AMÉDRO (2008, Fig. 14). This section spans several significant unconformities (Fig. 3B), one at the base of the Gault Clay overlying the Lower Greensand, a second at the condensed interval VIII zone, a third at the XII zone, and the highest at the Albian/Cenomanian contact between the Gault and the glauconitic marl (HART, 2000). The LOC of the lower interval of zones I-VII is poorly constrained at the top by the FO of *Actinoceramus sulcatus* and its slope is the same as that of the higher LOCs. The unconformity break is placed at the top of zone VIII. The LOC in the next higher interval of zones IX-XI is constrained by the range of *Callihoplites auritus*. In the next higher section of zones XII-XIII the LOC is constrained by several FO and LO bioevents (Fig. 3B). This section plot defines

the relative duration of the hiatus of two unconformities as meter thicknesses in the Mont Risou section; these intervals can now be projected into other graphed sections.

An older ammonite data set from Folkestone by AMÉDRO (1992) is graphed by several FO and LO's. The unconformities are well constrained as in the newer section in AMÉDRO (2008). The Merstham, England (AMÉDRO, 2008, Fig. 15) repeats the uppermost part of the Upper Greensand Formation that is present at Folkestone. The data are confined to two horizons so the LOC is not well constrained. However these data adjust nine FO's and nine LO's. The Strépy and Harches sections were graphed next (AMÉDRO, 2008, Fig. 17) and have limited data so that the LOC of each is not tightly constrained.

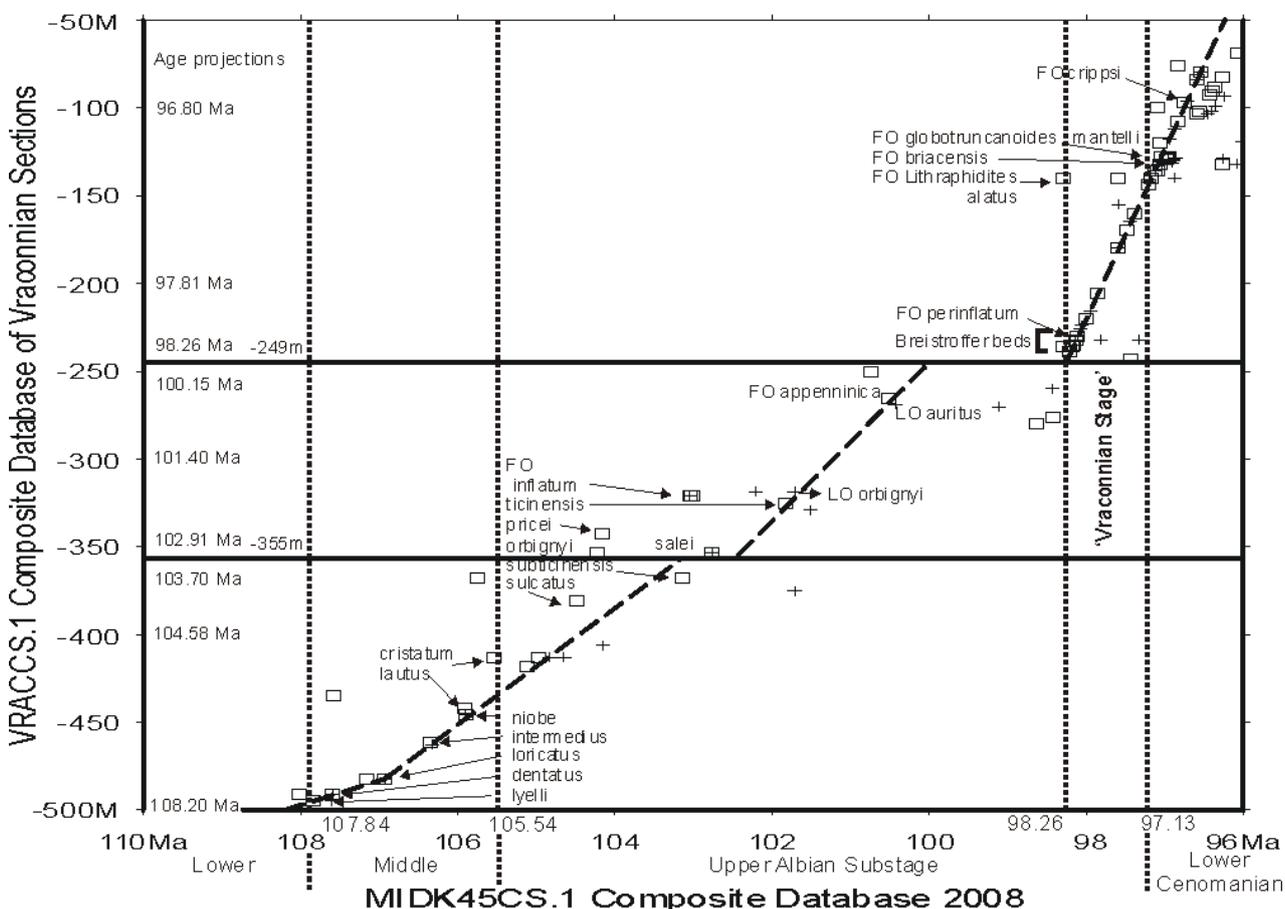


Figure 4: Graphic correlation plot of VRACCS.1 database to the global integrated MIDK45CS.1 database.

#### Process of calibration to numerical ages

The graphic correlation experiment resulted in a list of species and their ranges relative to each other in the metric scale of the Mont Risou section. The next step in the experiment was to convert the metric scale to a numerical age scale in mega-annum units. This was accomplished by plotting the composited range data set of AMÉDRO's 'Vraconnian' data set, VRACCS.1, to the MIDK45CS.1 composited range data set (Fig. 4). The MIDK45CS.1 data

set is the next development stage beyond MIDK3 (SCOTT *et alii*, 2000) and MIDK42CS.1 (SCOTT, in press; see also data of MIDK42 on website [precisionstratigraphy.com](http://precisionstratigraphy.com)). It is composed of more than one hundred sections and nearly 3000 bioevents, geochemical events, magnetochrons, and sequence stratigraphic contacts. The scale of this range data set is the numerical time scale of HARLAND *et alii* (1990), as revised by Ogg *et alii* (2004), with the exception of the age ascribed the base Cenomanian, 97.13 Ma, because correlations of

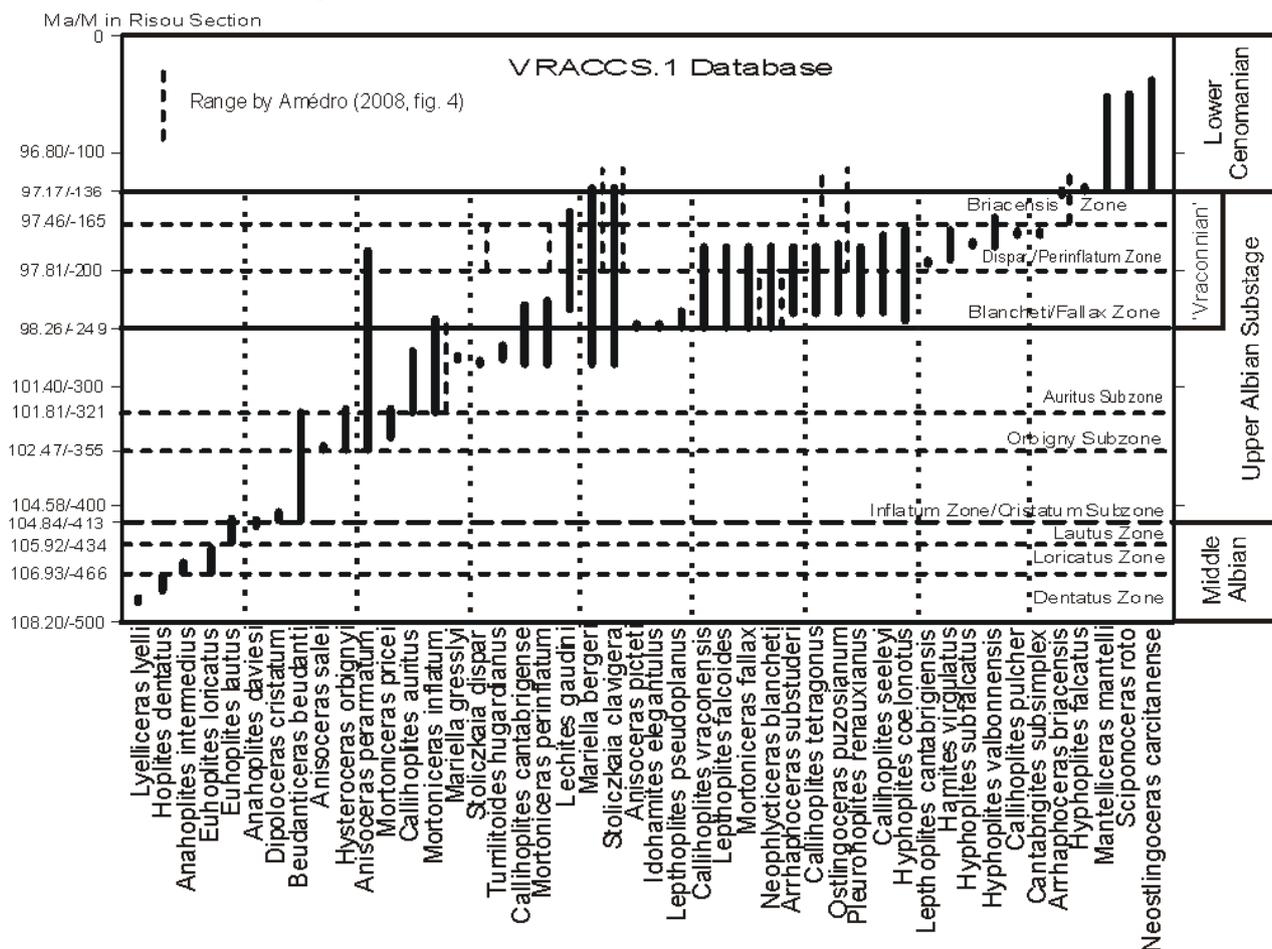
radiometrically dated bentonites and bioevents have been re-evaluated (OBOH-IKUENOBE *et alii*, 2007).

The two unconformities in the uppermost Albian Upper Greensand Formation are well defined by the data plot (Fig. 4). The two hiatuses separate the VRACCS.1 section into three intervals; the lower break is at -355 meters and the upper break is at -249 meters, which is the base of the 'Vraconnian Stage'. The duration of the lower hiatus is 103.70 to 102.91 Ma, and that of the upper hiatus 100.15 to 98.26 Ma. The LOC of the lower interval is well constrained by numerous bioevents. This lower interval represents two rates of sediment accumulation, a slower basal interval and a faster upper interval. This rate change is indicated by the change in the LOC slope. The LOC of the middle interval is tightly constrained by ammonites and planktic foraminifera. The upper interval includes numerous bioevents and the LOC is very tightly constrained. The age of the 'Vraconnian Stage' spans from 98.26 to 97.13 Ma. The black shale, organic-rich BREISTROFFER

marker beds were deposited at the beginning of the 'Vraconnian'.

**Integrated biostratigraphic ranges**

The stratigraphic ranges of 46 ammonites that were recorded in the seven key sections of AMÉDRO (2008) were integrated into a single range chart at the scale of the thickness of the Mont Risou section (Fig. 5). This scale was converted to a numerical age scale by graphing the composited VRACCS.1 data set to the MIDK45CS.1 data set as explained above. The three Middle Albian Substage zones are defined by the FO of the nominate species (Fig. 1 ; Table 2). The base of the Upper Albian Substage is defined by the FO of *Dipoloceras cristatum*, which is the basal subzone of the Inflatum Zone. Two of the next overlying subzones are defined in this dataset by the FOs of *Hysterocheras orbigny* and *Callihoplites auritus*; *Hysterocheras varicosum* is not included in this dataset. The base of *Stoliczkaia dispar* defines the base of the uppermost Upper Albian Dispar Zone.



**Figure 5:** Chronostratigraphic range chart of ammonites in 'Vraconnian' composited database of key 'Vraconnian' sections. Ages interpolated by graphic correlation of the VRACCS.1 database with the MIDK45CS.1 database. The ages of some ranges are younger in the subset of 'Vraconnian' sections than in the complete set of sections in the MIDK45CS database used in Figure 7.

| TAXA                          | VRACCS: FO - LO | MIDK45CS: FO - LO | Leptholites cantabrigiensis | -194.7543 | -194.7543                      | 97.75*    | 97.75*    |          |          |
|-------------------------------|-----------------|-------------------|-----------------------------|-----------|--------------------------------|-----------|-----------|----------|----------|
| Acanthoceras amphibolum       | 65.0472         | 65.0472           | 94.3454                     | 92.6212   | Leptholites falcoides          | -248.88   | -180.3551 | 98.58*   | 98.31*   |
| Acanthoceras rhotomagense     | 5.9825          | 43.151            | 94.9483                     | 92.5959   | Leptholites pseudoplanus       | -248.88   | -234.88   | 98.31*   | 98.16*   |
| Actinoceras subulcatus        | -406.00         | -406.00           | 104.79*                     | 104.79*   | Lithraphidites acutus          | -90.8027  | 101.6637  | 96.3969  | 92.7     |
| Actinoceras sulcatus          | -381.00         | -329.0148         | 104.4776                    | 101.5134  | Lithraphidites alatus          | -139.916  | -105.5228 | 98.296   | 93.2     |
| Algericeras boghariense       | -103.4988       | -93.7467          | 96.5874                     | 96.2374   | Lithraphidites carniolensis    | -236.00   | -80.00    | 134.1294 | 65.4901  |
| Algericeras proratum          | -92.6427        | -64.6744          | 96.4245                     | 95.578    | Lithraphidites pseudoquadratus | -236.00   | 21.0707   | 98.1749  | 93.87    |
| Amphizygus brooksii           | -236.00         | -80.00            | 124.717                     | 65.505    | Lyelliceras lyelli             | -475.7512 | -475.7512 | 107.84   | 107.5987 |
| Anahoplites daviesi           | -414.8517       | -410.6029         | 105.1156                    | 104.8296  | Manivitella permatoidea        | -236.00   | -80.00    | 133.9695 | 65.2514  |
| Anahoplites intermedius       | -465.8373       | -450.2584         | 107.1613                    | 106.32    | Mantelliceras dixoni           | -41.4901  | -41.49    | 95.6585  | 95.4     |
| Anisoceras perarmatum         | -353.00         | -180.0123         | 98.426                      | 97.5888   | Mantelliceras mantelli         | -132.00   | -38.00    | 97.0656  | 95.65    |
| Anisoceras picteti            | -248.88         | -247.2            | 94.8091                     | 94.3577   | Mantelliceras saxbii           | -127.8239 | -54.0022  | 96.9643  | 95.5     |
| Anisoceras pseudoelegans      | -188.2678       | -188.2678         | 97.68*                      | 97.68*    | Mariella bergeri               | -280.00   | -129.6378 | 98.6353  | 96.8887  |
| Anisoceras salei              | -353.00         | -353.00           | 102.7656                    | 102.7656  | Mariella cenomanensis          | -127.8239 | -112.3952 | 96.9524  | 95.7203  |
| Arrhaphoceras briacensis      | -136.00         | -132.00           | 97.0865                     | 97.0865   | Mariella gresslyi              | -276.00   | -276.00   | 98.59*   | 98.59*   |
| Arrhaphoceras substuderi      | -236.8768       | -180.3551         | 98.18*                      | 97.6*     | Markalius circumradiatus       | -228.00   | -80.00    | 134.0985 | 82.2933  |
| Axopodorhabdus albianus       | -236.00         | 102.7677          | 109.77                      | 90.6992   | Marker bed Breistroffer        | -235.00   | -224.00   | 98.1644  | 98.0493  |
| Axopodorhabdus dietzmannii    | -139.916        | 35.7909           | 128.6386                    | 82.56     | Mojsisoviczia subdelaruei      | -436.8038 | -436.8038 | 105.8965 | 105.8965 |
| Beudanticeras beudanti        | -410.6029       | -321.1158         | 104.9638                    | 103.0425  | Mortonoceras alstonensis       | -248.88   | -247.2    | 98.31*   | 98.29*   |
| Biticinella breggiensis       | -367.6759       | -250.0185         | 105.757                     | 97.5059   | Mortonoceras fallax            | -248.88   | -180.00   | 98.31*   | 97.58*   |
| Calculites anfractus          | -140.00         | -84.00            | 97.1702                     | 96.5842   | Mortonoceras inflatum          | -321.1158 | -241.59   | 103.0425 | 99.1149  |
| Callihoplites auritus         | -321.1924       | -269.043          | 103.0146                    | 100.4255  | Mortonoceras perinflatum       | -280.00   | -226.00   | 98.6353  | 98.0702  |
| Callihoplites cantabrigense   | -280.00         | -230.00           | 98.6353                     | 98.1121   | Mortonoceras pricei            | -342.7298 | -318.414  | 104.1574 | 102.1977 |
| Callihoplites leptus          | -241.587        | ***               | 98.23*                      | ***       | Neolobites vibrayanus          | 101.6637  | 102.7677  | 93.7514  | 93.12    |
| Callihoplites pulcher         | -170.00         | -170.00           | 97.49*                      | 97.49*    | Neophlycticeras blancheti      | -248.88   | -180.00   | 98.31*   | 97.58*   |
| Callihoplites seeleyi         | -236.00         | -170.00           | 98.17*                      | 97.49*    | Neostilgoceras carctanense     | -132.00   | -38.00    | 97.0656  | 96.1028  |
| Callihoplites tetragonus      | -236.8768       | -180.00           | 98.18*                      | 97.60*    | Ostlingoceras puzosianum       | -236.00   | -129.6378 | 98.17*   | 96.97*   |
| Callihoplites vracconensis    | -248.88         | -180.0123         | 97.49*                      | 97.60*    | Oxytyridoceras roissyanum      | -472.9187 | -472.9187 | 108.0147 | 107.58   |
| Cantabrigites subsimplex      | -170.00         | -170.00           | 97.49*                      | 97.49*    | Percivalia haxtonensis         | -144.00   | -144.00   | 97.2121  | 93.47    |
| Carbon peak OAE 2             | 100.9276        | 104.9757          | 93.52                       | 93.45     | Placozygus fibuliformis        | -230.00   | -80.00    | 98.1121  | 65.505   |
| Chiastozygus bifarius         | -236.00         | -84.00            | 98.1749                     | 65.505    | Planomalina buxtorfi           | -236.00   | -119.00   | 100.606  | 96.0093  |
| Chiastozygus litterarius      | -236.00         | -80.00            | 128.2573                    | 65.5535   | Planomalina praebuxtorfi       | -250.0185 | -216.0556 | 100.7341 | 97.9467  |
| Chiastozygus platyrhethus     | -236.00         | -80.00            | 125.5054                    | 88.3575   | Pleurohoplites renauxianus     | -236.8768 | -180.3551 | 98.18*   | 97.60*   |
| Corollithion kennedyi         | -80.00          | 101.6637          | 96.55                       | 92.00     | Pleurohoplites subvarians      | -180.0123 | -180.0123 | 97.59*   | 97.59*   |
| Corollithion madagaskarensis  | -236.00         | -80.00            | 122.358                     | 65.3262   | Praeglobotruncana delrioensis  | -222.9565 | 101.6637  | 108.0343 | 92.00    |
| Corollithion signum           | -236.00         | -80.00            | 114.0455                    | 70.4437   | Praeglobotruncana stephani     | -235.7826 | 118.2239  | 100.509  | 90.41    |
| Cretarhabdus conicus          | -236.00         | -80.00            | 133.1464                    | 64.5992   | Prediscosphaera columnata      | -236.00   | -80.00    | 122.8533 | 90.7     |
| Cretarhabdus striatus         | -236.00         | -80.00            | 117.1136                    | 65.505    | Prediscosphaera cretacea       | -84.00    | -80.00    | 119.3508 | 64.523   |
| Cribrosphaerella ehrenbergii  | -236.00         | -80.00            | 116.0163                    | 94.3685   | Prediscosphaera spinosa        | -220.00   | -80.00    | 121.9656 | 64.4953  |
| Dicarinella algeriana         | 38.73           | 118.2239          | 95.105                      | 89.7362   | Pseudaspidoceras flexuosum     | 109.7598  | 109.7598  | 93.044   | 93.00    |
| Dicarinella hagni             | 80.3194         | 118.2239          | 93.8363                     | 89.1675   | Quadrans gartneri              | 107.1837  | 128.16    | 93.5451  | 68.3833  |
| Dimorpholites niebe           | -449.5502       | -436.8038         | 106.3502                    | 105.8965  | Rhagodiscus achlyostaurion     | -236.00   | -80.00    | 124.6741 | 92.00    |
| Dipoloceras cristatum         | -413.00         | -406.00           | 105.5386                    | 104.1574  | Rhagodiscus angustus           | -236.00   | -80.00    | 123.278  | 65.505   |
| Discorhabdus ignotus          | -236.00         | -80.00            | 128.6386                    | 65.505    | Rhagodiscus asper              | -236.00   | -80.00    | 128.6386 | 92.95    |
| Eiffellithus turrisseiffelii  | -236.00         | -80.00            | 101.856                     | 64.4399   | Rhagodiscus splendens          | -228.00   | -80.00    | 123.278  | 65.551   |
| Ellipsagelospaera ovata       | -216.00         | -112.00           | 121.959                     | 88.1021   | Rotalipora appenninica         | -265.787  | 23.6467   | 100.5159 | 94.48    |
| Eprolithus apertior           | -236.00         | -80.00            | 121.1235                    | 96.5423   | Rotalipora brotzeni            | -136.00   | -119.00   | 97.0842  | 94.62    |
| Eprolithus floralis           | -236.00         | -84.00            | 122.8468                    | 85.5995   | Rotalipora cushmani            | 33.9508   | 101.6637  | 96.17    | 93.07    |
| Euhoplites lautus             | -433.9713       | -410.6            | 105.9188                    | 104.6453  | Rotalipora deeckeii            | 23.6467   | 37.9989   | 95.3597  | 93.2     |
| Euhoplites loricaeus          | -465.8373       | -436.8038         | 106.9309                    | 105.8965  | Rotalipora evoluta             | -139.916  | -52.1622  | 97.588   | 95.05    |
| Euhysterochoceras nicaisi     | -103.4988       | -19.04            | 96.5874                     | 94.5395   | Rotalipora gandolfi            | -222.9565 | 88.7835   | 98.9078  | 93.52    |
| Euomphaloceras septemseriatum | 108.6557        | 108.6557          | 93.88                       | 93.15     | Rotalipora globotruncanoides   | -136.00   | 101.6637  | 97.13    | 93.15    |
| Favusella washitensis         | -139.916        | 11.8706           | 114.1975                    | 94.84     | Rotalipora greenhornensis      | -100.0028 | 99.8236   | 97.0842  | 93.07    |
| Forbesiceras beaumontianum    | -127.8239       | -128.8911         | 96.9524                     | 96.2621   | Rotalipora montsalvensis       | -59.5223  | 101.6637  | 97.384   | 93.15    |
| Gartnerago nanum              | -236.00         | -96.00            | 98.296                      | 94.23     | Rotalipora reicheli            | -1.0096   | 32.1108   | 96.706   | 94.42    |
| Gartnerago obliquum           | 107.1837        | 128.16            | 99.2536                     | 64.743    | Rotalipora crenulata           | -236.00   | -80.00    | 124.612  | 65.597   |
| Gartnerago praeobliquum       | -84.00          | ***               | 96.5842                     | ***       | Rotalipora laffitei            | -224.00   | -80.00    | 134.0402 | 66.407   |
| Gartnerago stenostaurion      | -139.916        | -128.3391         | 97.588                      | 96.864    | Scaphites simplex              | -172.3889 | ***       | 97.51*   | ***      |
| Gartnerago theta              | -108.00         | -80.00            | 96.8353                     | 89.0722   | Schloenbachia varians          | -119.9188 | -38.00    | 97.0648  | 95.12    |
| Graysonites azregensis        | -127.82         | -127.2379         | 96.95*                      | 96.94*    | Sciponoceras roto              | -132.00   | -50.00    | 97.0656  | 96.2284  |
| Graysonites cobbiani          | -126.0738       | -122.2903         | 96.92*                      | 96.87*    | Sharpeiceras laticlavicum      | -76.0825  | -42.2261  | 96.8389  | 95.45    |
| Hamites virgulatus            | -191.1179       | -165.9805         | 97.71*                      | 97.45*    | Sharpeiceras schlueteri        | -127.8239 | -103.4988 | 96.9524  | 96.4478  |
| Hayesites albiensis           | -232.00         | ***               | 118.6477                    | 97.3296   | Staurolithes glaber            | -232.00   | -112.00   | 98.133   | 96.8772  |
| Hedbergella libyca            | -236.00         | -212.00           | 100.6217                    | 94.66     | Stoliczkaia africana           | -222.9565 | -131.2738 | 98.04*   | 97.00*   |
| Helenea chiastia              | -224.00         | 102.7677          | 133.9695                    | 92.7      | Stoliczkaia clavigera          | -280.00   | -132.00   | 98.6353  | 97.0656  |
| Helicolithus trabeculatus     | -236.00         | -80.00            | 122.8533                    | 65.505    | Stoliczkaia dispar             | -280.00   | -280.00   | 98.6353  | 97.116   |
| Hoplites dentatus             | -472.9187       | -465.8373         | 107.5987                    | 106.93    | Stoverius achylosus            | -172.00   | -80.00    | 123.3079 | 97.116   |
| Hyphoplites coelonotus        | -242.942        | -165.00           | 97.4319                     | 97.4319   | Tegumentum stradneri           | -230.00   | -80.00    | 129.3543 | 83.3875  |
| Hyphoplites falcatus          | -132.0523       | -132.0523         | 96.2621                     | 96.08     | Tetrapodorhabdus coptensis     | -230.00   | -80.00    | 128.4545 | 93.05    |
| Hyphoplites subfalcatus       | -179.0295       | -179.0295         | 97.51*                      | 97.51*    | Ticinella praeticinensis       | -367.6759 | -325.2222 | 110.9318 | 97.4606  |
| Hyphoplites valbonnensis      | -180.00         | -155.0337         | 97.5888                     | 97.5888   | Ticinella primula              | -393.1481 | -270.6389 | 111.614  | 97.3667  |
| Hypoturrilites gravesianus    | -128.8911       | -112.3309         | 96.9684                     | 94.84     | Ticinella subticinensis        | -367.6759 | -232.00   | 103.1399 | 97.82    |
| Hypoturrilites schneegansi    | -127.8239       | -98.9             | 96.9524                     | 96.3536   | Ticinella ticinensis           | -325.2222 | -140.00   | 101.8279 | 96.8779  |
| Hysterochoceras orbigny       | -353.00         | -318.414          | 104.226                     | 101.71    | Tranolithus gabalus            | -230.00   | -80.00    | 123.9877 | 83.5584  |
| Idohamites elegantulus        | -248.88         | -247.2            | 98.31*                      | 98.29*    | Tranolithus minimus            | -220.00   | -132.00   | 98.0074  | 65.505   |
| Inoceramus anglicus           | -130.00         | -96.00            | 104.1574                    | 96.7098   | Tranolithus orionatus          | -230.00   | -80.00    | 110.9137 | 69.1743  |
| Inoceramus concentricus       | -435.00         | -375.0571         | 107.5807                    | 101.71    | Turrilites acutus              | 26.5908   | 71.6713   | 94.64    | 93.98    |
| Inoceramus crispus            | -97.00          | -38.00            | 96.7471                     | 94.2      | Turrilites costatus            | -23.8259  | 26.5908   | 95.3943  | 94.14    |
| Lechites gaudini              | -236.00         | -151.2174         | 98.6353                     | 96.9133   | Turrilites scheuchzerianus     | -14.8097  | -0.6416   | 96.1343  | 94.07    |
| Lechites moreti               | -276.00         | -276.00           | 98.59*                      | 98.59*    | Turrilitoides hugardianus      | -276.00   | -260.00   | 98.426   | 98.426   |

**Table 2:** Biostratigraphic ranges of Albian Stage ammonites, planktic foraminifera and nannofossil zones in 'Vraconnian' sections in metric units in the Mont Risou section (VRACCS.1) and numerical ages in the global MIDK45CS.1 database; ages with asterisk are interpolated by plotting the 'Vraconnian' database, VRACCS.1 to the MIDK45CS.1 database.

AMÉDRO (2008) proposed that the base of the 'Vraconnian Stage' be defined by the FOs of *Mortonicerias fallax* and *Neophlycticeras blancheti*. He included three ammonite zones in this stage in ascending order: *Mortonicerias (Mortonicerias) fallax* Interval Zone, *Mortonicerias (Subschloenbachia) perinflatum* Total Range Zone, and the *Arrhaphoceras (Praeschloenbachia) briacensis* Interval Zone (Fig. 1). The composited dataset of his sections shows that *M. inflatum* first appears lower in the section than *M. fallax* (Fig. 5) because GALE *et alii* (1996) reported it lower in their Mont Risou section than did AMÉDRO (2008, Fig. 11). The FO of *A. briacensis* is at the base of the Cenomanian Stage based on this set of sections. The ranges of AMÉDRO's key zonal species (AMÉDRO, 2008, Fig. 4) are dashed lines on Fig. 5 ; some ranges are similar and others are quite different from the ranges derived by graphic correlation of his data. *S. dispar* has a short range because it occurs in two sections, the Diégo well and

GALE's *et alii* Mont Risou section. Both *Mariella bergeri* and *Stoliczkaia clavigera* are low in the Mont Risou section (GALE *et alii*, 1996). Some species have longer ranges in AMÉDRO's dataset because they occur with species of other zones in condensed intervals in other sections not graphed, such as at the 'Vraconnian' stratotype (RENZ & JUNG, 1978) and near Drap in the Alpes-Maritimes, France (DELANOY & LATIL, 1988).

This graphic correlation experiment shows that the base of the *S. dispar* Zone and the FO of *M. perinflatum* are significantly lower (31 m) and older (380 kyr) than the FO of either *M. fallax* or *N. blancheti*. Thus the concept of a 'Vraconnian Stage' is not equal to the Dispar Zone. However the FOs of *M. perinflatum*, *M. fallax/N. blancheti*, and *A. briacensis* may be used to divide the Dispar Zone into three subzones.

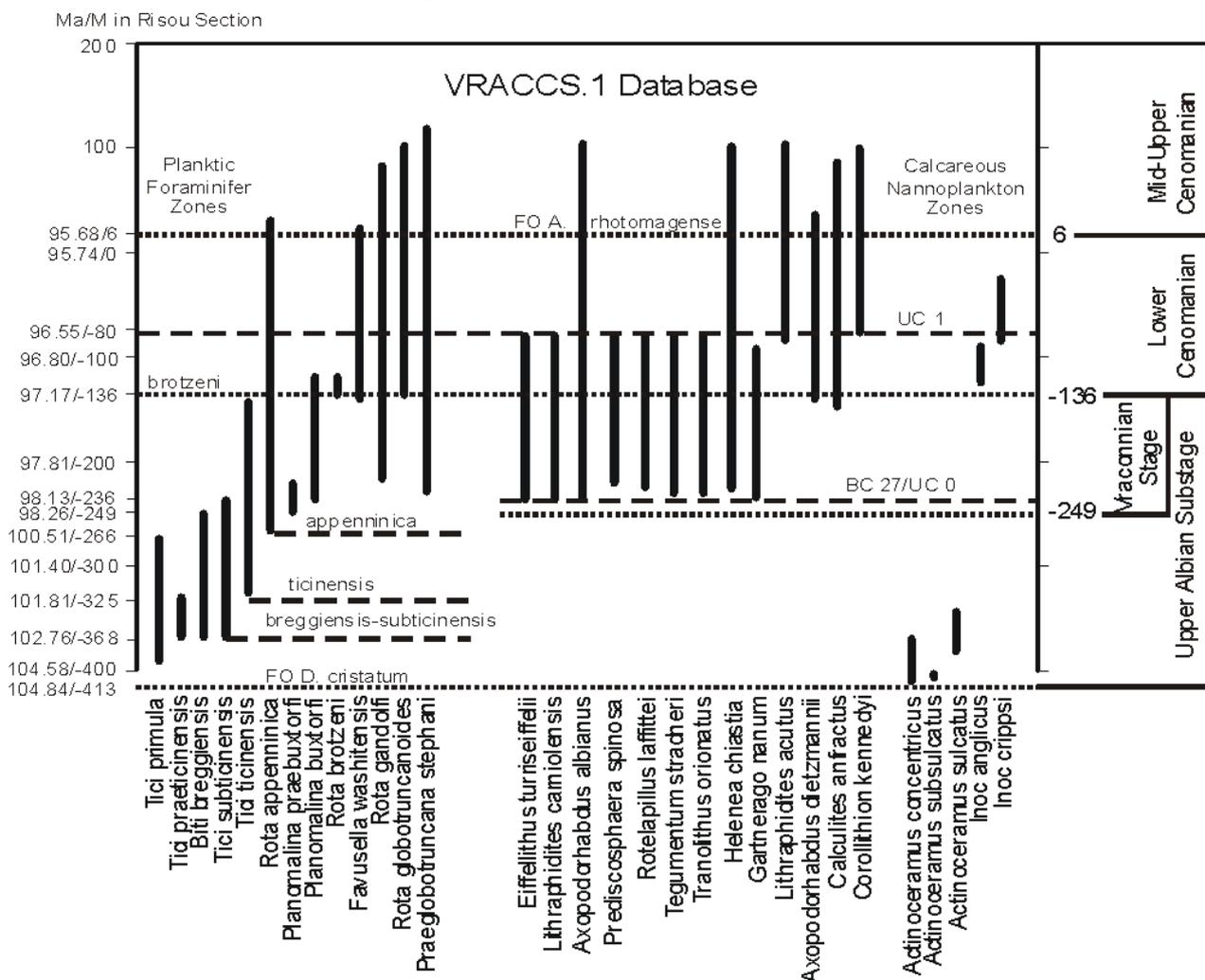


Figure 6: Chronostratigraphic range chart of planktic foraminifera, calcareous nannofossils, and inoceramids in 'Vraconnian' composited database of key 'Vraconnian' sections. Vertical axis is meters in the Mont Risou section. Ages in Ma are interpolated by graphic correlation of the VRACCS.1 database with the MIDK45CS.1 database. Highest part of composite section not converted to ages. Foraminifer zones are defined by FO of named taxa. Nannofossil zones are defined by FO of *E. turrisieffeli* and *C. kennedyi* respectively. The FO of *Lithraphidites acutus* is in the middle part of the range of the Lower Cenomanian ammonite *Mantelliceras mantelli* in the Kalaat Senan section, Tunisia and 103.5 m above the base of *Rotalipora globotruncanoides* (MIDK.10 section; ROBASZYNSKI *et alii*, 1994).

Integration of key planktic foraminifera and calcareous nannofossils (Fig. 6) based on the limited dataset of AMÉDRO (2008) shows that *R. appenninica* is below the base of *Mortonicerax fallax*, and the first occurrence of *E. turriseiffelii* is slightly above it. In the larger MIDK45 dataset the FO of *E. turriseiffelii* is projected at 101.86 Ma and *R. appenninica* at 100.37 Ma, both of which are significantly older than the FO of *M. fallax* at 98.26 Ma. So neither pelagic species is useful as a proxy guide to the base of a 'Vraconnian Stage.'

Five inoceramid species are recorded in the sections at Mont Risou and at Folkestone (Fig. 6). *Actinoceramus concentricus*, *Actinoceramus sulcatus*, and *Actinoceramus subsulcatus*

characterize the lower part of the Upper Albian and *Inoceramus anglicus* and *Inoceramus cripsii* characterize the Lower Cenomanian.

The Albian stage is divided into seven zones and twenty-five subzones. The ranges of key fossils that define these zones can be calibrated to numerical Ma ages by graphic plots to sections bearing dated bentonites and geochemical events. This process measures the durations of the zones as proposed by AMÉDRO and ROBASZYNSKI (2008). This database is anchored to bentonites in the U.S. Western Interior dated by OBRADOVICH (1993) and projected to the age of Magnetochron M0 at the base of the Aptian. As new radiometric ages are accrued this database can be tested and adjusted to accommodate new data.

|                          |                         |                         |                           |                     |                     |        |                       |                          |        |
|--------------------------|-------------------------|-------------------------|---------------------------|---------------------|---------------------|--------|-----------------------|--------------------------|--------|
| Albian                   | Upper                   |                         | <i>S. (F.) blancheti</i>  | 98.26*              | <i>M. fallax</i>    | 98.26* | <i>R. appenninica</i> |                          |        |
|                          |                         | M. inflatum             | <i>C. auritus</i>         | 103.01              | <i>M. inflatum</i>  | 98.64  | 100.52                |                          |        |
|                          |                         |                         | <i>H. varicosum</i>       | 103.03              | M. pricei           | 104.16 | <i>R. ticinensis</i>  | <i>E. turriseiffelii</i> |        |
|                          |                         |                         | <i>H. orbigny</i>         | 104.23              |                     |        | 101.83                | 101.91                   |        |
|                          |                         |                         | <i>D. cristatum</i>       | 105.54              | <i>D. cristatum</i> | 105.54 | <i>B. breggiensis</i> | <i>E. monechiae</i>      |        |
|                          |                         |                         |                           |                     | 105.76              | 101.97 |                       |                          |        |
|                          | Middle                  | E. lautus 106.23        | <i>A. daviesi</i>         | 105.68              |                     |        |                       |                          |        |
|                          |                         | E. loricatus 107.16     | <i>E. nitidus</i>         | 105.78              |                     |        |                       |                          |        |
|                          |                         |                         | <i>E. meandrinus</i>      | NA                  |                     |        |                       |                          |        |
|                          |                         |                         | <i>M. subdelaruei</i>     | 105.9               |                     |        |                       |                          |        |
|                          |                         |                         | <i>D. ricobe</i>          | 106.35              |                     |        |                       |                          |        |
|                          |                         | H. (H.) dentatus 107.60 | <i>A. intermedius</i>     | 107.16              |                     |        |                       |                          |        |
|                          |                         |                         | <i>H. (H.) spathi</i>     | 107.6               |                     |        |                       |                          |        |
|                          |                         |                         | <i>L. lyellii</i>         | 107.84              |                     |        |                       |                          |        |
|                          | <i>L. pseudolyellii</i> |                         | NA                        |                     |                     |        |                       |                          |        |
|                          | Lower                   | D. mammillatum 111.58   | <i>P. (L.) steinmanni</i> | 108.2               |                     |        |                       |                          |        |
|                          |                         |                         | <i>O. bulliensis</i>      | 109.47              |                     |        |                       |                          |        |
|                          |                         |                         | <i>P. puzosianus</i>      | 110.8               |                     |        |                       |                          |        |
|                          |                         |                         | <i>O. raulinianus</i>     | 109.39              |                     |        |                       |                          |        |
|                          |                         |                         | <i>C. floridum</i>        | 111.31              |                     |        |                       |                          |        |
|                          |                         |                         | <i>S. kitcheni</i>        | NA                  |                     |        |                       |                          |        |
|                          |                         |                         | <i>D. perifrata</i>       | NA                  |                     |        |                       |                          |        |
|                          |                         | L. tardefurcata         | <i>L. (N.) regularis</i>  | 110.94              |                     |        |                       |                          |        |
|                          |                         |                         | <i>L. tardefurcata</i>    | 112.66              |                     |        |                       |                          |        |
|                          |                         |                         | <i>L. schrammeni</i>      | NA                  |                     |        |                       |                          |        |
| <i>T. praeticinensis</i> |                         |                         | 110.93                    | <i>T. orionatus</i> |                     |        |                       |                          |        |
| <i>T. primula</i>        |                         |                         | 111.61                    | 110.91              |                     |        |                       |                          |        |
|                          |                         | <i>H. planispira</i>    | 129.19                    | <i>C. signum</i>    |                     |        |                       |                          | 114.05 |

**Figure 7:** Chronostratigraphic chart of the original Albian Stage. Ammonite zones from OGG *et alii* (2004), HOEDEMAEKER *et alii* (2003), HOEDEMAEKER & RAWSON (2000), HOEDEMAEKER *et alii* (1993), HANCOCK (1991), and OWEN (1984a, 1984b). Planktic foraminifera zones based on FOs from PREMOLI SILVA & SLITER (2002). Nannofossil zones based on FOs from BOWN *et alii* (1998) and BRALOWER *et alii* (1995). Numerical ages interpolated by graphic correlation of MIDK45 database (SCOTT *et alii*, 2000). NA indicates species not in the MIDK45 database.

### Chronostratigraphic correlation of key sections

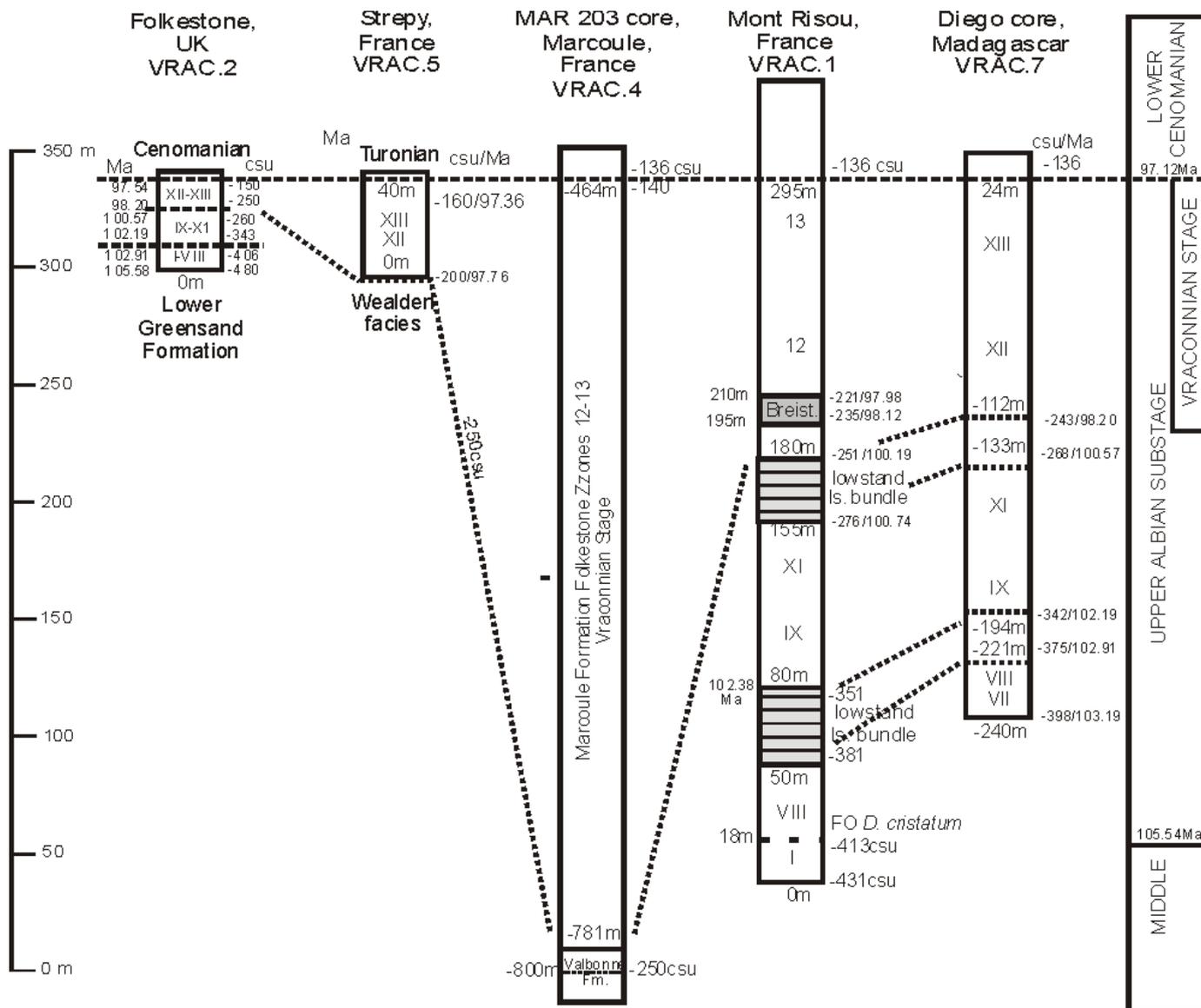
The graphic correlation method provides data for chronostratigraphic correlation of the key 'Vraconnian' sections (Fig. 8). The proposed 'Vraconnian Stage' correlates with the upper part of the Upper Albian Substage (Fig. 4). The cross section datum is the Albian/Cenomanian stage boundary as defined by the FO of *R. globotruncanoides*. AMÉDRO used the FOs of *Mortonicerax fallax* and *Neophlycticeras blancheti* to define the base of the 'Vraconnian', which is approximated by the -250 m position

in the VRACCS.1 database. This position correlates with the unconformity at the base of zones XII-XIII in the Folkestone section and with the transgressive facies between -800 and -781 m in the MAR 203 core at Marcoule. The same time line also correlates within the low-stand limestone bundles in the Mont Risou and Diégo core. The entire interval of the Bracquignies Formation in the Strépy boatlift section and in the Harchies N° 1 well (AMÉDRO, 2008, Fig. 17) correlates with Folkestone zones XII-XIII. The organic-rich BREISTROFFER interval in the Mont Risou section lies within the lower part of the transgressive interval equivalent to zone XII; it may represent maximum flooding. In the

Mont Risou section a lower bundle of bioclastic-glaucouitic limestone from about 50 to 80 m (AMÉDRO, 2008, Fig. 11) correlates with the lower condensed section between zones VIII and IX at Folkestone.

The thickness of the 'Vraconnian Stage' varies greatly among the sections as noted by (AMÉDRO, 2008). In the MAR 203 core at Marcoule the equivalent 'Vraconnian' interval is 317 m thick, eight times than thicker than at the Strépy section, more than twenty-three times thicker than at Folkestone, and nearly

160 times thicker than in the 'Vraconnian' reference section in Switzerland. The rates of sediment accumulation varied from 4.5 m/myr at Folkestone to 37.6 m/myr at Mont Risou, and 111.6 m/myr at Marcoule. The rate of accumulation is based on the compacted section and is not a sedimentation rate. These rates are based on the duration of about 3 myr for the 'Vraconnian' interval. This great range in rates is based on very different basin subsidence histories and tectonic conditions of each section.



**Figure 8:** Stratigraphic correlation of key 'Vraconnian' sections using base of Cenomanian Stage as datum. CSU values are meters in the Mont Risou section of GALE *et alii* (1996, MIDK.24). Ages interpolated by graphic correlation of the VRACCS.1 database with the MIDK45CS.1 database. Age of lower part of Valbonne Formation below intraformational unconformity at -800 m not projected because of the absence of fossils.

## Re-examination of the rationale for a 'Vraconnian Stage'

### 1. The interval is mappable

This criterion is essential to the definition of lithostratigraphic units such as formations (NACSN, 2005). However, it is not part of the definition of a stage, in which lithofacies change from basin to basin. The lithologies that comprise the 'Vraconnian' interval are quite different from section to section (AMÉDRO, 2008), and thus do not make up a mappable lithostratigraphic unit.

### 2. The interval records an important eustatic event

Indeed the 'Vraconnian' interval records a third-order three myr depositional cycle of transgression and regression on a regional even global scale. This feature, however, defines sequence stratigraphic units, not stages (NACSN, 2005). The Upper Albian Stage records five such sequences of this scale (SCOTT *et alii*, 2003), but it would be impractical to divide the Albian into five stages.

### 3. The interval has a distinctive and diverse fossil assemblage

This property is an essential feature of a stage concept. Three species of *Mortoniceras* and one species of *Stoliczkaia* comprise the 'Vraconnian Stage' according to AMÉDRO (2008) and AMÉDRO & ROBASZYNSKI (2008). AMÉDRO (2008) characterizes the 'Vraconnian' by the abrupt diversification of heteromorph ammonites of the Turrilitidae, Hamitidae, Anisoceratidae, and Baculitidae in the condensed La Vraconne section, where they comprise 60% of the specimens. However, species of three of these families appear earlier in the Late Albian between 102.47 and 98.64 Ma (Fig. 5) earlier than the FO of *Mortoniceras fallax* and *Neophlycticeras blancheti*.

The three heteromorph species restricted to the 'Vraconnian' in AMÉDRO's database (2008) are primarily found in Western Europe. In the Carpatho-Balkan region of Eastern Europe the 'Vraconnian' is represented by a condensed interval of glauconitic and phosphatic sandy limestone less than one meter thick; the only ammonite species of the zone is *Stoliczkaia notha* (KUTEK & MARCINOWSKI, 1996). One North American section, Dry Creek in northern California (AMÉDRO, 2008, Fig. 24), yields three ammonite species of the Dispar Zone, only one of which, *Lechites gaudini*, is found at the type section of the 'Vraconnian' in Switzerland (RENZ & JUNG, 1978). Few species characteristic of the 'Vraconnian' are found outside of Western Europe. The ammonite assemblage is not widespread in the Tethyan or Boreal Realms.

### 4. The interval duration is similar to that of other stages

The proposed duration of the 'Vraconnian Stage' is 2-3 myr, which is equivalent to that of the Santonian Stage. However the durations of Cretaceous stages vary from 2.3 to 13 myr (OGG *et alii*, 2004) and duration is not a criterion for defining a stage.

### 5. The interval locally is quite thick

Sections bearing the 'Vraconnian' fauna are locally thicker than the underlying part of the Albian. The thickness difference is highly variable from basin to basin and within basins. Such thickness differences are found between condensed sections and coeval basin margin and basin center sections of many zones.

## Conclusions

The diverse ammonite assemblage in the uppermost part of the Upper Albian Substage comprises a distinctive zone in Western Europe that can be subdivided into three subzones. These subzones are recognized in many Tethyan and transitional Boreal sections in Europe. However, at the 'Vraconnian' type section in Switzerland *Stoliczkaia dispar* and *Mortoniceras perinflatum* are the only zonal named taxa present. The strata bearing the *S. dispar* Zone are bounded by unconformities in many sections and they are but one of five Upper Albian depositional sequences. The thickness of this sequence varies from condensed sections of two meters to expanded basinal sections more than 300 meters thick. The *S. dispar* Zone represents a time interval of about three myr, which is about the same duration as the briefest Cretaceous ages. This interval is very useful as a biostratigraphic unit and a third-order sequence stratigraphic unit. However this interval is not a practical chronostratigraphic unit such as a stage because its boundaries cannot be demonstrated to be synchronous and the interval is not isochronous, nor is it globally recognizable. Defining this interval as a stage equivalent to the Albian and Cenomanian stages would materially alter the concept of the Albian Stage by deleting its uppermost zone. The concept of a 'Vraconnian Stage' is not a practical subdivision of the Cretaceous System.

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