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# CONODONT SUCCESSION ACROSS THE TOURNAISIAN-VISEAN BOUNDARY BEDS AT SALET, BELGIUM

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KEY WORDS : Conodonta, Tournaisian, Visean, Boundary, Salet, Dinant Synclinorium.

#### INTRODUCTION.

The taxonomy of the conodont genera as *Siphonodella*, *Gnathodus*, *Pseudopolygnathus*, *Scaliognathus*, *Mestognathus*, that exhibit a great importance for the Lower Carboniferous stratigraphy have recently been revised by SANDBERG *et al.* (1978), LANE, SANBERG, ZIEGLER (1980), Helka (1983) and LANE & ZIEGLER (1983). New concept of several species as well as descriptions of new taxa stimulated us to re-investigate the conodonts occurring in the classical sections of the Dinant Synclinorium. The purpose of this study is also to attempt an application, in the Dinantian type area, of the preliminary standard conodont zonation proposed by LANE, SANDBERG & ZIEGLER (1980) for the post-*Siphonodella* interval. This study offers a revision of the conodont fauna recovered from the section called "Route de Salet" which is exposed along the Molignée Valley, in a small quarry and it continues along the road to Salet (Text-figs 1-2). The precise location and accessibility of this exposure is given by CONIL & GROESSENS (1974 : 24), GROESSENS (1975 : 45) and HANCE (1985). It is a classical and one of the most important sections for the Lower Carboniferous stratigraphy. As a parastratotype section for the Lower Viséan (CONIL, 1967) it attracted the interest of many workers being the area of an intense search for microfossils (see HANCE, 1985). Several foraminiferal taxa (*cf.* CONIL & LYS, 1964) have been



Fig. 1 - General location map. Patterned area shows extent of the Carboniferous deposits in the Dinant Synclinorium.

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described from this exposure for the first time and it is also a type locality for two stratigraphically important conodont species, *Dollymae bouckaerti* GROESSENS and *Eotaphrus bultyncki* (GROESSENS). The holotytypes of these conodonts are reillustrated herein.

The section at Salet (Textfig. 2), moreover, is a type section for two lithostratigraphical units of the classic Dinantian of Belgium, the Calcaire Noir de la Malignée (GROESSENS, 1975) and the Calcaire de Salet (GROESSENS, CONIL & LEES, 1976). The base of the Moliniacian, i.e. the stage of the Viséan, locally distinguished in Belgium, was also defined here at the Bed 52 (PAPROTH *et al.*, 1983 : 188).

The investigated conodont fauna includes the material collected and originally described by GROESSENS (1971, 1975) and also conodonts obtained during additional recent sampling that produced some new conodont occurrences. The samples were collected from the lower portion of the sequence only up to the Bed 285 where there occur slump structures at the top of the Calcaire de la Molignée (Text-fig. 2).

#### LITHOLOGY.

At Salet, a 320 meter thick section of the Lower Carboniferous carbonates is exposed, but only 130 m were covered by this study. The investigated portion of the section consists of three lithological units (Text-fig, 2) that constitute proximal lateral equivalents of the Waulsortian facies (*cf.* LEES, 1984).

The lowest unit is developed as crinoidal and bryozoan packstones that are massive or interbedded with thin shales in the lower part of this unit. These sediments pass laterally into the Bayard facies considered to be located in the neighborhood of the Waulsortian buildups.

Overlying is the Leffe facies, which comprises well-bedded wackestones and packstones that contain various allochems and first of all peloids, bioclasts of echinoderms and bryozoans, and coated grains. The Leffe facies extends landwards for many kilometers away from the buildups (LEES & CONIL, 1980; LEES, 1982) and it always separates the latter from the Molignée faciès.

The Calcaire de la Molignée forms the upper unit sampled for conodonts in the "Route de Salet" section. The limestones are wackestones and lime mudstones with numerous shaly interbeds. The composition of allochems is similar to that of the Leffe facies, but from the Bed 87 the increase of foraminifers in place of bryozoans is noted to occur. According to HANCE (1985 : 182), the lower part of the Calcaire de la Molignée at Salet reflects regressive conditions, whereas the Viséan transgression is indicated much later by the first appearance of the foraminiferal fauna dominated by the Archaediscidae in the Bed 215.

#### CONODONT DISTRIBUTION,

Very closely sampling of long intervals in the "Route de Salet" section provided substantial account of conodont succession in the late Tournaisian and early Viséan strata. Although the observed occurrences of the majority of conodont species (Table 1) agree with their recognized stratigraphic ranges (*cf.* LANE, SANDBERG & ZIEGLER, 1980), some species, however, demonstrate new occurrences that appear to be very significant for the phylogeny of these species and for the Lower Carboniferous conodont stratigraphy.

The most remarkable is the first appearance of *Gnathodus texanus*, a zonal marker of the *texanus* Zone. The lower range of this species was known to start simultaneously with the extinction of *Scaliognathus anchoralis* (LANE, SANDBERG & ZIEGLER, 1980 : Tab. 2; LANE & ZIEGLER, 1983). In contrast, *G. texanus* occurs at Salet just below the appearance of *Scaliognathus anchoralis europensis* (Text-fig. 3). Similary to that, the ranges of both species are recently noted to overlap in the Dinantian rocks of southwestern Ireland (THORNBURY, 1985) and VARKER & SEVASTOPULO (1985 : Tab. 6) indicated that for the Lower Carboniferous condonts in Britain and Ireland the range of *Gnathodus texanus* extends from the middle



Fig. 2 - The lower part of the "Route de Salet" section sampled for conodonts. Black triangle indicates the position of the Tournaisian-Viséan boundary. Bed numeration after Overlau et Conil, 1965.

part of the *anchoralis* Zone. The record of *G. texanus* at Salet does not appear to be a freak of nature. It is well documented in our fauna by transitional forms (see Table 1 and P1. 6, Figs 7-9) showing the evolutionary relation between *Gnathodus* semiglaber and Gnathodus texanus. The former species gave rise to G. texanus by the reduction of the inner parapet and platform ornamentation. The reduction trend of morphological changes in the conodont platform, as late Tournaisian gnathodontan Pa elements display (BELKA 1984 : fig. 10; 1985b) is also exemplified by the origin of Gnathodus pseudosemiglaber from G. cuneiforms. This is indicated by transi-tional forms (Pl. 5, Figs. 4-9) as well as by ontogenetic development of small specimens of G. pseudosemiglaber which recapitulate a character of *G. cuneiformis*. To show phyletic sequence of above mentioned species of *Gnathodus*, the transitional forms are separately presented in Table 1, but they are not considered herein to be the separate species.

The other anomaly which is worthy to note, is the record of *Pseudopolygnathus pinnatus*. This species (P1. 2, Fig. 9) yields earlier appearance relative to *Pseudopolygnathus oxypageus*, *Dollymae bouekaerti*, and *Eotaphrus bultyneki*. It is much earlier than was recognized by LANE, SANDBERG & ZIEGLER (1980 : Tab. 2) but identical range of *Ps. pinnatus* reflecting, most probably, an actual stratigraphic range of this species is recorded from Britain and Ireland (VARKER & SEVASTOPULO 1985 : Tab. 6).

The genus Scaliognathus is represented by Sc. praeanchoralis (Pl. 1, Fig. 7), Sc. anchoralis fairchildi (Pl. 1, Figs. 8-11) and Sc. anchoralis europensis (Pl. 2, Fig. 2). LANE & ZIEGLER (1983 : 205-206) suggested that Sc. a. europensis probably developed from Sc. a. fairchildi although the appearance of Sc. a. fairchildi before Sc. a. europensis was not documented at that time. Now, such an evidence is recorded. In the "Route de Salet" section Sc. a. fairchildi comes 1.5 m before Sc. a. europensis.

The species Gnathodus homopunctatus, which together with Mestognathus beckmanni is considered to be diagnostic for earliest Viséan strata (GROESSENS 1975), appears much later (Text-fig. 3) than in other sections of the Dinant Basin. At the Tournaisian-Viséan boundary stratotype in Dinant it occurs before *M. beckmanni* following the Bed 141, at the base of which the boundary is placed (GROESSENS & NOEL, 1977 : Pls 1-2). New data, however, suggest that the range of *G. homopunctatus* may extend down to the middle of the *anchoralis* Zone, as formerly recorded in the Moravia-Silesia Basin in Poland (BELKA, 1985 : Fig. 3).

#### CORRELATION.

It was already the preliminary investigation of conodonts in several sections of the Dinantian of Belgium (GROESSENS, 1971) that showed these microfossils as excellent guides for correlation of the Tournaisian rocks. As a result, moreover, prospects of recognition of conodont zones in the Tn3a-V1b interval has also been outlined at that time. These zones were incorporated later by AUSTIN (1973) into his proposal of the Dinantian conodont zonation suggested for application in Europe.

The conodont zonal scheme for the Lower Carboniferous currently used in Belgium was established by GROESSENS (1975). Recently, it was in part modif-ied to produce some new detailed sub-divisions (PAPROTH *et al.*, 1983 : Tab. Although the usefulness of this 2). scheme for the Belgian stratigraphy remains unquestionable, its global application proves to be limited. This is because some species used for definition of subzones (e. g. Dollymae hassi, Dollymae bouckaerti, Eotaphrus bultyncki) are rather rare out of Belgium while other index forms as Polygnathus communis carinus and Eotaphrus burlingtonensis appear in Belgium significantly later than their lower ranges recognized in many Lower Carboni-ferous sections (see LANE, SANDBERG & ZIEGLER, 1980). Undoubtedly, this is why the framework of the world-wide pre-liminary standard conodont zonation (SANDBERG *et al.*, 1978; LANE, SANDBERG & ZIEGLER, 1980) differs considerably from that recognized in Belgium by GROESSENS (1975).

The investigated section yields all diagnostic forms which allowed to identify all the conodont zones of Upper Tournaisian and Lower Viséan strata sensu GROESSENS (1975), i. e. the carinus Zone, the anchoralis Zone, and the beckmanni Zone (Text-fig. 3), and also to correlate this section with other successions of Belgium (GROESSENS, 1975; GROESSENS, CONIL & LEES, 1976).

Of special interest in this classical section, however, was also to test up the preliminary standard conodont zonation. The results we received show the scheme is to be corrected.

The typicus Zone is recognized and its base is placed at the lowest occurrence of Gnathodus cuneiformis (Bed 22/69), the lower range of which appears to be very close to that of Gnathodus typicus Morphotype 2. In the investigated section, the zonal name-bearer appears erratically but much later than G. cuneiforms (see Table 1). Besides all that the zone is well documented by associated fauna including other gnathodids, as well as Pseudopolygnathus pinnatus, Ps. oxypageus, Dollymae bouckaerti, and Eotaphrus bultyncki.

In fact, the base of the typicus Zone can be identified in Belgium as well as in other regions of Europe by the first appearance of *G. cuneiformis*. Thus, it is only proper to replace the typicus Zone by the *cuneiformis* Zone, as proposed by BELKA (1985 : 46-47). The prominence of *Gnathodus typicus* in the studies of LANE, SANDBERG & ZIEGLER (1980) is not reflected within the composition of the Lower Carboniferous conodont faunas from Europe. Moreover, this species can be also difficult to recognize (THORNBURY, 1985 : 35-36). As a result of the examination of conodont fauna from Salet we



Fig. 3 - Ranges of the most important condont taxa around the Tournaisian-Viséan boundary beds in the "Route de Salet" section M. - Mestognathus, Sc. a. - Scaliognathus anchoralis, Pr. - Protognathodus, Ps. - Pseudopolygnathus. The recognized standard condont zones (on the right) are compared with those (on the left) of GROESSENS (1975) and CONIL, GROESSENS & PIRLET (1977). Lithology : 1 - massive limestones, 2 - bedded limestones, 3 - bedded limestones with cherts, 4 - shales; for detailed lithology, see HANCE (1985). expect that further studies will allow to subdivide the *cuneiformis* Zone into two or three parts. The records of easily recognizable forms as *Protognathodus cordiformis*, *Pseudopolygnathus pinnatus* and/or *Dollymae bouckaerti* seem to be very perspective.

Since the species Scaliognathus anchoralis has been subdivided to include three subspecies, the base of the anchoralis Zone is defined by the first appearance of Sc. a. europensis (LANE, SANDBERG & ZIEGLER, 1980; LANE & ZIEGLER, 1983 : Fig. 4). The zonal marker succeeds at Salet Sc. a. fairchildi (Table 1). Therefore, the base of the anchoralis Zone must be taken 1.5 m higher than it was traced out formerly (GROESSENS, 1975 : 46). The new concept of this limit is more useful in comparison to that of GROESSENS (1975), if there are disparities in lower ranges of particular subspecies of Scaliognathus anchoralis. Moreover, it provides a mutual control of this boundary by the occurrences of other forms of the genus, such as Sc. praeanchoralis and Sc. a. fairchildi.

The top of the anchoralis Zone being the base of the next texanus Zone is identified by the first appearance of Gnathodus texanus (LANE, SANDBERG & ZIEGLER, 1980 : 120). As mentioned above, this taxon appears in the investigated section earlier than Sc. a. europensis (Table 1 and Text-fig. 3). Consequently, the anchoralis Zone would be to eliminate here the if following the concept of LANE, SANDBERG & ZIEGLER. This zone, however, constitu-tes the best recognized interval of the Lower Carboniferous that by means of conodonts can be traced throughout the world. Therefore, it is suggested to re-examine the concept of the texanus Zone in order to create another zone of a global use that will follow the *anchoralis* Zone. long as the stratigraphic ranges of other Viséan gnathodids are not satisfactorily documented, Mestognathus beckmanni can be only tentatively used to define this zone. The species Mestognathus beckmanni is interpreted to have lived in shallow-water biofacies (AUSTIN, 1976; BELKA, 1983 : 81-82) whereas the preliminary standard condont zonation for the Lower Carbonife-rous is essentially based on deep-water conodonts.

# THE POSITION OF THE TOURNAISIAN-VISEAN BOUNDARY.

The Tournaisian-Viséan boundary as ratified by the Subcommission on Carboniferous Stratigraphy runs 34 meters below the base of the Marbre Noir de Dinant and it is located precisely at the base of Bed 141 of the boundary stratotype section at Dinant (*cf.* CONIL *et al.*, 1969). LANE & ZIEGLER (1983) discussed recently in detail the conodont occurrences around the boundary in that section. As a consequence,they correlated the Tournaisian-Viséan boundary with the preliminary standard conodont zonation to put it within the *anchoralis* Zone.

In the "Route de Salet" section, the Tournaisian-Viséan boundary has been distinguished at the base of Bed 8/52 (Text-figs 2 and 3) that forms the base of the Marbre Noir de la Molignée. The

recognition is based, first of all, on such lithological evidences as the presence of an important argillaceous marker (*cf*.CONIL, GROESSENS & PIRLET, 1977) that occurs in the first meter above the boundary. In terms of conodonts and foraminifers the present position of the limit appears to be njustified. The order of ranges of conodonts such as Scaliognathus anchoralis europensis and Mestognathus beckmanni (Text-fig. 3) in relation to the appearance of the diagnostic foraminifer *Pachysphaerina pachysphaerica* is identical to that in the Tournaisian-Viséan Boundary Stratotype at Dinant (*cf.* GROESSENS & NOEL, 1977 : Pl. 1). The taxon *Scaliognathus anchoralis europensis* (together with Hindeodella segaformis), namely, ranges up to Bed 8/54 (Table 1) i.e. 0.5 m higher than hitherto was recognized, while Mestognathus beckmanni starts in Bed 8/60. The foraminifer Pachysphaerina pachysphaerica the first appearance of which is coincident at Dinant with the Tournaisian-Viséan bound-ary, ranges from Bed 8/57 (HANCE, 1985 : Tab. 18).

Although the range of *Mestognathus* beckmanni in the stratotype section at Dinant is considered to be artificial (LANE & ZIEGLER, 1983 : 210) the biostratigraphic data evidence that the Tournaisian-Viséan boundary at Salet should be displaced higher up to the base of Bed 8/57 (see Text-fig. 3).

#### SYSTEMATIC PALEONTOLOGY.

All type material and figured specimens are housed in the collection of the Geological Survey of Belgium, Brussels.

Genus Staurognathus BRANSON & MEHL, 1941.

TYPE SPECIES : Staurognathus cruciformis BRANSON & MEHL, 1941.

#### REVISED DIAGNOSIS :

A genus characterized by an Icriodus-like P element with a long anterior process bearing a double row of side denticles which may be fused to form transverse ridges. As many as two lateral processes and one shorter posterior process may be developed to produce the typical cruciform outline. Lateral processes are straight or very slightly concave anteriorly. They may also be ornamented by a double row of denticles on the upper surface. The posterior process is generally curved in wardly.

#### REMARKS :

The genus Staurognathus, as presently conceived, comprises two species : St. cruciformis BRANSON & MEHL, and St. dionantensis n. sp. which is described herein. The another earlier described species, Staurognathus anchorarius HASS (1959) has been transferred by CHAUFF & KLAPPER (1978) to Bactrognathus as it bears only a single row of denticles on the anterior process and it shows a morphologic gradation with B. excavatus. The species St. dionantensis appears to be an ancestor of St. cruciformis, at it is evidenced by a transitional form described by AUSTIN & GROESSENS (1972) as "New genus B GROESSENS  $\rightarrow$  Staurognathus BRANSON & MEHL".

The both species of Staurognathus yield distinct exclusive geographic restriction in their distribution. The type species has so far been found to occur widely in North America (BRANSON & MEHL, 1941; COOPER, 1948; HASS, 1959; BURTON, 1964; THOMPSON, 1967; THOMPSON & FELLOWS, 1970; LANE, SANDBERG & ZIEGLER, 1980; CHAUFF, 1981). In contrast, St. dionantensis as a very rare element was only reported from Belgium, Germany (cf. LANE, SANDBERG & ZIEGLER, 1980), and Poland (BELKA, 1985a). The single Australian specimen of Staurognathus cruciformis (DRUCE, 1970), represents, most probably, another species of the genus just as forms described by CHAUFF (1985 : Figs. 1.27 and 1.29) as Staurognathus aff. S. cruciformis.

The hypothetic apparatus reconstruction of *Staurognathus cruciformis* was presented by CHAUFF (1981). Except for Pa and Pb elements this apparatus is believed to be identical with that of the genus *Doliognathus*. CHAUFF (1985 : 308) suggested, moreover, that *St. cruciformis* have evolved from an early form of *D. latus*.

The morphological similarities between St. cruciformis and some forms of D. latus, as demonstrated by CHAUFF (1985), are unquestionable. However, St. cruciformis with well-developed lateral processes is known to occur stratigraphically below the first appearance of Doliognathus latus (see LANE, SANDBERG & ZIEGLER, 1980; Tab. 2; CHAUFF, 1981 : Text-fig. 3 and Tab. 2). We believe that the evolution of D. latus to develop an inner lateral process reflects the general evolutionary trend observed among the uppermost Tournaisian conodonts to produce symmetrical Pa elements. This trend is easily recognizable in the evolution of the genus Scaliognathus for instance (cf. LANE & ZIEGLER, 1983).

RANGE :

Base of the *cuneiformis* Zone into the *anchoralis* Zone.

#### Staurognathus dionantensis n. sp. Pl. 1, Figs 1-2

- 1959 Icriodus latericrescens BRANSON & MEHL ? -VOGES, p. 286 (specimens not illustrated).
- 1971 N. GEN. B. GROESSENS, p. 17, P1. 2, Figs. 5-6.
- 1980 Eotaphrus ? n. sp. V. LANE, SANDBERG & ZIEGLER, Pl. 10, Figs. 9-10 (specimens from VOGES' collection).
- 1985 Eotaphrus ? sp. V of LANE, SANDBERG & ZIEGLER. BELKA, p. 37, Pl. 9, Fig. 6.

HOLOTYPE :

The specimen illustrated by GROESSENS (1971, Pl. 2, Fig. 5a-b) and reillustrated herein in Pl. 1, Fig. 2.

DERIVATIO NOMINIS :

From Dinant, where this species was found for the first time in Belgium.

#### STRATUM TYPICUM :

Bed of crinoidal packstone, Bed 8/0 of the Route de Salet, 20 m below the base of the Marbre Noir de la Molignée.

LOCUS TYPICUS :

Salet, Molignée Valley, the Ardennes; Belgium.

DIAGNOSIS : A species of *Staurognathus* characterized by only germinal development of lateral processes. The posterior process is marked by a group of nodes or a short ridge that diverges (at the angle of 45°) inwards from the axis of the anterior process. The large open basal cavity flares beyond the spindle. The posterior margin of the basal cavity is corrugated to form three lobe-like extensions.

MATERIAL : 5 sepcimens.

REMARKS :

Since the time when this element has been recognized it is difficult to assign it, beyond any doubt, to any Lower Carboniferous conodont genus. This was because it displays many similarities, but also differences to such genera as *Icriodus, Eotaphrus, Staurognathus* and *Dollymae* (*cf.* GROESSENS, 1976). Recently, this unit was commonly regarded as a possible species of *Eotaphrus*, but lack of a large cusp, diagnostic for *Eotaphrus*, was noticed. Although a resemblance to the genus *Eotaphrus* was suggested, a very close connection with the genus *Staurognathus* was also indicated (AUSTIN & GROESSENS, 1972).

LANE, SANDBERG & ZIEGLER (1980) confined the stratigraphic range of this species to the lower part of the anchoralis Zone. The appearance of this species earlier than Scaliognathus anchoralis was, however, known in Belgium (GROESSENS, CONIL & LEES, 1976). Recently, in the Moravia-Silesia Basin in Poland, a single specimen of this species was found to occur at the base of the cuneiformis Zone (BELKA, 1985a). This record precedes considerably the lower limit of Eotaphrus evae, which is considered to be the most primitive form of Eotaphrus. Thereby, a phylogenetic relation of the newly described species to the Eotaphrus lineage (as given by LANE, SANDBERG & ZIEGLER 1980) seems to be doubtful. As a consequence, we assign this species to Staurognathus, on the basis of the tendency in morphologic gradation to form the cruciform outline.

The new species, Staurognathus dionantensis n. sp. differs from St. cruciformis in lacking of well-developed posterior and lateral processes. It resembles Eotaphrus burlingtonensis, which can be, however, distinguished by the prominent cusp at the posterior end.

The origin of *St. dionantensis* is problematic. Two forms, at first *Peleksygnathus* sp. A of VOGES and later *Dolymae* sp. A of VOGES are suggested to have been an ancestor of this species (*cf.* AUSTIN & GROESSENS, 1972; GROESSENS, 1976).

RANGE : Same as for the genus.

	Zone				unzoned					interval							unzoned interval																						
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TABLE I - Numerical data on the conodont collection from the "Route de Salet" section. Conodont sample numbers are bed numbers.

	Zone				cuneiformis								cuneiformis													a	ncho	rali	s									
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G. punctatus			-	-	-				2 1	-		-		-	-	-	-	-	-		-		-	-		-				-	-	-	-				-	-
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## PLATE CAPTIONS

### PLATE I

All upper view, except as noted.

- Figs. 1-2 Staurognathus dionantensis n. sp. 1 - Bed 8/0, x65 2 - Holotype, Bed 8/0, x45.
- Eotaphrus bultyncki (GROESSENS, 1971) Rephotograph of holotype (GROESSENS, 1971; Pl. 1, Figs. 4a, b), Fig. 3 Bed 22/92, x75, lateral view.
- Figs. 4-6 Dollymae bouckaerti GROESSENS, 1971 4 - Rephotograph of holotype (GROESSENS, 1971; Pl. 1, Figs. 7a, b)

  - Kepholograph of horocype (interesting)
    Bed 8/0, x55.
    5 Bed 8/0, x60, lower view.
    6 Bed 8/0, x90, Juvenile specimen.
- Scaliognathus praeanchoralis LANE, SANDBERG & ZIEGLER, 1980 Bed 8/3, x100. Fig. 7

Figs.8-11 Scaliognathus anchoralis fairchildi LANE & ZIEGLER, 1983 8a, b - Bed 8/1C, x75, lower and upper views. Specimen with larger posteriorly directed cusp. 9-Bed 8/2, x115. Juvenile specimen.

- 10 Bed 8/3, x130. Juvenile specimen. 11 Bed 8/1A, x85. Specimen transitional to Scaliognathus anchoralis europensis.



Fig.	1	Doliognathus dubius BRANSON & MEHL, 1941. Bed 8/2, x100, upper view. Juvenile specimen.
Fig.	2	Scaliognathus anchoralis europensis LANE & ZIEGLER, 1983. Bed 8/51, x70, upper view.
Figs	.3a,b	Mestognathus beckmanni BISCHOFF, 1957 Bed 8/67, x50, upper and lateral views.
Fig.	4	Bispathodus bispathodus ZIEGLER, SANDBERG & AUSTIN, 1974. Bed 8/57, x85, upper view.
Figs	5a,b	"Spathognathodus" laterigranosus GEDIK, 1974. Bed 8/2, x95, lateral and upper views.
Figs	6-7	"Spathognathodus" macer (BRANSON & MEHL, 1934). 6 - Bed 8/17, x65, lateral view. 7a,b - Bed 8/56, x60, upper and lateral views.
Fig.	8 -	Anchignathodus simplicatus (RHODES, AUSTIN & DRUCE, 1969). Bed 8/68, x150, lateral view. Juvenile specimen.
Fig.	9 -	Pseudopolygnathus pinnatus (VOGES, 1959). Bed 8/1B, x40, upper view.
Figs.	10-12	<ul> <li>2 - Pseudopolygnathus multistriatus MEHL &amp; THOMAS, 1947.</li> <li>10 - Bed 22/86, x40, upper view. Morphotype 1.</li> <li>11 - Bed 22/92, x45, upper view. Morphotype 2.</li> <li>12 - Bed 22/92, x50, upper view, Morphotype 2.</li> </ul>

Fig. 13 - Pseudopolygnathus oxypageus LANE, SANBERG & ZIEGLER, 1980. Bed 8/3, x65, upper view. Morphotype 2.

Contraction of the

plate 2



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Figs. 1 - 2	Polygnathus cf. purus VOGES, 1959. 1 - Bed 22/86, x90, upper view. 2 - Bed 22/86, x95, lower view.
Figs. 3 - 4	Polygnathus communis communis BRANSON & MEHL, 1934. 3 - Bed 8/57, x75, upper view. 4 - Bed 8/57, x105, lower view. Juvenile specimen.
Fig. 5	Polygnathus flabellus (BRANSON & MEHL, 1938). Bed 8/51, x60, upper view.
Figs. 6 - 7	<i>Polygnathus bischoffi</i> RHODES, AUSTIN & DRUCE, 1969. 6 - Bed 8/44, x75, upper view. 7 - Bed 8/44, x50, lower view. Advanced form.
Figs. 8 - 10	Polygnathus inornatus E. R. BRANSON, 1934. 8 – Bed 8/158, x75, upper view. Specimen transitional to P. flabellus.
	9 - Bed 8/56, x70, upper view. 10a, b - Bed 8/44, x50, lower and upper views.
Figs. 11 - 15	<ul> <li>Polygnathus longiposticus BRANSON &amp; MEHL, 1934.</li> <li>11 - Bed 8/0, x75, lower view. Juvenile specimen with a wide, inverted basal cavity. Compare with large specimen (P1. 3, Fig. 14a) showing an extremely small basal cavity.</li> </ul>
	12 - Bed 8/0, x50, lower view.
	13 - Bed 8/3, x60, lower view.
	14a,b - Bed 8/17, x40, lower and upper views. Large, advanced form with a generatic lobe-like extension of the platform margin
	15 - Bed 8/17, x40, upper view.

plate 3



	All upper views, except Fig. 5.
Fig. 1	Protognathodus praedelicatus LANE, SANDBERG & ZIEGLER, 1980. Bed 22/86, x75.
Figs. 2 - 5	<ul> <li>Protognathodus cordiformis LANE, SANDBERG &amp; ZIEGLER, 1980.</li> <li>2 - Bed 8/3, x85. Juvenile specimen.</li> <li>3 - Bed 8/3, x75.</li> <li>4 - Bed 8/1C, x85. Juvenile specimen.</li> <li>5 - Bed 8/2, x85. Margin of the cup partially broken.</li> </ul>
Fig. 6	Gnathodus punctatus COOPER, 1939. Bed 22/86, x70.
Figs. 7 -10	Gnathodus delicatus BRANSON & MEHL, 1938. 7 - Bed 8/17, x60. Specimen transitional to G. punctatus. 8 - Bed 8/1A, x85. 9 - Bed 8/0, x75. 10 - Bed 22/69, x75.
Figs. 11-12	Gnathodus typicus COOPER, 1938, Morphotype 1. 11 - Bed 8/0, x70. 12 - Bed 8/0, x55.
Figs. 13-16	Gnathodus cuneiformis MEHL & THOMAS, 1947. 13 - Bed 22/88, x45. 14 - Bed 8/0, x90. 15 - Bed 8/53, x75. Younger morphotype. 16 - Bed 8/1A, x80.

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PLATE 4



Figs. 1 - 3 Gnathodus cuneiformis MEHL & THOMAS, 1947.
1 - Bed 8/98, x70.
2 - Bed 8/1A, x55. Specimen transitional to G. delicatus.
3 - Bed 8/1A, x80. Specimen with a peculiar transverse ridge perpendicular to the blade.
Figs. 4 - 9 Gnathodus cuneiformis MEHL & THOMAS, 1947 → Gnathodus pseudosemiglaber THOMPSON & FELLOW, 1970.
4 - Bed 8/53, x95. Juvenile specimen.
5 - Bed 8/7, x80.
6 - Bed 8/97, x85. Juvenile specimen.
7 - Bed 8/104, x60.
9 - Bed 8/104, x55.

Figs. 10 - 15 Gnathodus pseudosemiglaber THOMPSON & FELLOWS, 1970.

10 - Bed 8/104, x50.
11 - Bed 8/104, x50.
12 - Bed 8/104, x55.

Figs. 10 - 15 Bed 8/76, x75.

14 - Bed 8/91, x60.
15 - Bed 8/98, x60.

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All upper views.

Figs.	1 -	6	Gnathodus 1 - Bed 2 - Bed 3 - Bed 4 - Bed 5 - Bed 6 - Bed	semiglaber (BISCHOFF, 1957). 8/1C, x75. 8/1C, x75. 8/1C, x60. 8/1C, x75. 8/44, x40. 8/1A, x60.
Figs.	7 -	9	Gnathodus Gnathodus 7 - Bed 8 - Bed 9 - Bed	<pre>semiglaber (BISCHOFF, 1957) -&gt; texanus ROUNDY, 1926. 8/1C, x75. 8/1A, x95. Juvenile specimen. 8/1C, x70.</pre>
Figs.	10 -	11	<i>Gnathodus</i> 10 - Bed 11 - Bed	texanus ROUNDY, 1926. 8/1C, x90. Juvenile specimen. 8/76, x65.
Figs.	12 -	14	Gnathodus 12 - Bed 13 - Bed 14 - Bed	praebilineatus BELKA, 1985. 8/163, x45. 8/190, x45. 8/190, x100. Juvenile specimen.

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Figs.	1 - 3	Gnathodus pseudosemiglaber THOMPSON & FELLOWS, 1970. 1 - Bed 8/122, x55. 2 - Bed 8/260, x75. 3 - Bed 8/104, x60.
Figs.	4 - 5	Gnathodus pseudosemiglaber THOMPSON & FELLOWS, 1970 → Gnathodus girtyi HASS, 1953. 4 - Bed 8/268, x70. 5 - Bed 8/263, x95.
Figs.	6 - 7	Gnathodus symmutatus RHODES, AUSTIN & DRUCE, 1969. 6 - Bed 8/83, x80. 7 - Bed 8/211, x120. Juvenile specimen.
Figs.	8 - 10	Gnathodus cf. homopunctatus ZIEGLER, 1962. 8 - Bed 8/263, x102. 9 - Bed 8/216, x 90. 10 - Bed 8/205, x 80.
Figs.	11 - 15	<ul> <li>Gnathodus homopunctatus ZIEGLER, 1962.</li> <li>11 - Bed 8/211, x140. Juvenile specimen.</li> <li>12 - Bed 8/211, x155. Juvenile specimen with an asymmetrical ornamented cup.</li> <li>13 - Bed 8/211, x135. Juvenile specimen.</li> <li>14 - Bed 8/178, x110. Specimen transitional to G. mermaidus.</li> <li>15 - Bed 8/183, x 95.</li> </ul>



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