Post-Pleistocene Sea Level Changes in the Baie Verte Estuary (Northumberland Strait) as Revealed by Foraminiferal Studies

by

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ABSTRACT

Two cores were obtained from the Baie Verte estuary of the Northumberland Strait to study relative sea level changes by examination of foraminiferal distributions.

The base of the cores contained a layer of salt marsh peat which can be accurately used to relocate former sea levels. Distinct vertical faunal assemblages observed in the cores indicate that there has been a change in the foraminiferal environments of the estuary during the last 5,000 years. The development of the estuary has resulted in the original, nontransitional estuary being transformed into the transitional system which exists today in the area. Evidence suggests that this change was due to alteration of circulation patterns in the estuary.

The implication for sea level changes indicates that a short period of emergence occurred at approximately 5,000 years B.P. This was followed by a continuous submergence to the present time. Thus, the evidence provides support for the models which have been developed to explain the migration of the peripheral bulge that followed deglaciation in the Maritimes.

A comparison of this study with one done in the same area by McRoberts (1968) indicates variations in the results obtained. In particular, large populations of arenaceous foraminifera found in the cores of the present study were not observed in McRoberts' (1968) core. The

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result has been different interpretations of events in the area related to relative sea level changes since the last glaciation.

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CHAPTER I

INTRODUCTION

General:

In recent years, a substantial amount of scientific research has been focused on the usefulness of foraminiferal distributions as sea level indicators in the Maritime Provinces (Scott, 1977; Scott and Medioli, 1978a, 1980). In particular, the attention was concentrated on using the information obtained from such studies to reconstruct an outline of the relative sea level changes that have occurred since the last glaciation (McRoberts, 1968; Scott, 1977; Scott and Medioli, 1978b, 1979). As a part of continuing effort towards establishing detailed relative sea level records in different parts of the Maritimes, many areas have been sampled. One of the most important of these, a small area in the Northumberland Strait, is the subject of this thesis.

The complex pattern of relative sea level rise and fall observed in the Maritimes has been attributed to the deformation of the lithosphere caused by the migration of a "peripheral bulge" that formed in the marginal regions of the ice sheet during the last glaciation (e.g. Peltier and Andrews, 1976; Scott, 1977). The phenomenon of migration of the peripheral bulge has been quantified in several

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mathematical models (Peltier and Andrews, 1976; Quinlan, 1978). All such models, however, are based on assumptions on the size of the last Laurentide ice sheet, the elasticity of the lithosphere and viscoelasticity of the asthenosphere. The only way to calibrate the assumptions is that of making predictions by means of the model and verifying them in the field. If the predictions are wrong, then the assumptions must be modified. Hence, there is the need for accurate and detailed measurements in the field.

The application of these models to the Maritimes, supported by existing data, suggests that the western part of the Northumberland Strait is now on the distal end of the "peripheral bulge" (which is migrating westward) and consequently, the relative sea level is steadily rising (Scott, 1977). The specific area of interest in the present study is the Baie Verte estuary, bordering on New Brunswick and Nova Scotia and located in this western portion of the Northumberland Strait.

Scope of Work:

Previous coring in the Baie Verte area (Kranck, 1972) had revealed the presence of peat layers but sea level results from these cores were questionable (McRoberts, 1968). Since the old cores no longer existed, it was necessary to return to the area where new cores were obtained (Fig.1,Table 1). It was hoped that foraminiferal assemblage zones could be



Figure 1. Core Locations in Baie Verte

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identified in the cores and related to modern estuarine and marsh environments. These results would help to establish the paleoecology of the area during this specific period of sedimentation. This information could be used to interpret the relative sea level changes in the development of this estuary. The sea level changes could then be related to the general studies on the migration of the "peripheral bulge" following deglaction.

More specifically, there was much interest to determine if the peat layers reported from previous cores (Kranck, 1972; McRoberts, 1968) were of freshwater or salt marsh origin. If the latter was the case, then extremely accurate sea levels could be derived using marsh foraminifera as indicators (Scott and Medioli, 1978a, 1980). It should be noted here that the organic material found in the cores of the present study is presently being C-14 dated. These dates were unavailable for incorporation into this study.

It was felt that one of these cores should be taken from the exact location as one obtained and analyzed by McRoberts (1968). In this way, the two could be compared in terms of foraminiferal distributions.

CHAPTER II

PREVIOUS WORK

Post-glacial History and Sea Level Investigations:

McRoberts (1968) conducted a study in which she examined the post-glacial history of the Northumberland Strait based upon distributions of foraminifera in two cores obtained by Kranck (1972). One of McRoberts' (1968) cores was from the same location as one examined in this study. She found a peat layer at the base of this core, describing it as depauperate of foraminifera and very rich in fresh water diatoms. She interpreted this as indicating fluvial deposits. Above this, an oyster shell layer was encountered. Samples examined by McRoberts above the basal peat layer consisted of a prolific Cribrononion fauna (=Elphidium in McRoberts, 1968). Arenaceous foraminifera were relatively rare in her samples. The upper section of the core was also recorded as depauperate and McRoberts concluded that this represented conditions similar to those existing in the modern Northumberland Strait.

In a study by Grant (1970) regarding recent costal My we have a submergence in the Maritime Provinces, he suggests that many areas in the Maritimes are still experiencing excessive submergence compared to the eustatic rate. Grant attributes this to a combination of crustal deflection by water load

from post-glacial eustatic rise of sea level and relaxation of the "peripheral bulge".

An extensive examination of the geomorphological development and post-Pleistocene sea level changes in the Northumberland Strait was carried out by Kranck (1972). Included in this study were two cores taken from the Baie Verte estuary at the same depths as the cores in this thesis (Table 1). The suggestion was made that "the Northumberland Strait may be on the edge of a region undergoing subsidence to compensate for post-glacial rebound in the centre of the continent (e.g. Newman and March, 1968). The area may straddle a fulcrum point separating a region of positive crustal movement from a region of negative crustal movement". (Kranck, 1972, p. 844)

Based on foraminiferal distributions in the cores taken from the estuarine - marsh systems at Chezzetcook, Chebogue, and Wallace, Nova Scotia, Scott (1977) observed differential relative sea level rises. He concluded that this could be attributed to the migration of the "peripheral bulge" following deglaciation in the Maritimes.

Additional studies are presently being conducted in the Maritimes by Scott and Medioli (1978b, 1979) to provide a more complete picture of relative sea level changes.

Foraminiferal Studies Related to This Work:

To obtain more precision in measuring former relative sea-levels, a method has been developed using vertical

zonations of marsh foraminifera as accurate indicators (Scott, 1977; Scott and Medioli, 1978a, 1980).

A study of the Miramichi River estuary was carried out by Scott <u>et al</u>. (1977). This study demonstrated that the distribution of foraminifera could provide a good indicator of circulation patterns and could be used to identify estuaries as either transitional or non-transitional. It was also shown by Scott <u>et al</u>. (1977) that faunal changes in estuaries can occur in a short time due to altered tidal circulation patterns.

Foraminifera and Environmental Factors:

It should be noted here that only species observed in the cores from this study will be discussed below.

Studies of the marsh foraminifera by Scott and Medioli (1972a, 1980) suggest that arenaceous species such as <u>Trochammina inflata and Tiphotrocha comprimata</u> occur only in the upper half of the salt marsh; this suggests that they are tolerant of long periods of exposure. Hence, since normal marshes inhabit only the upper half of the tidal range (Chapman, 1960), the presence of these arenaceous marsh forms limits a marine deposit to the upper quarter of the tidal range.

It has been suggested by Barlett (1966) that the proliferation of calcareous foraminifera in lower estuarine areas (in cool temperate climates) can be attributed to relatively higher salinities. He demonstrated that most

calcareous species such as <u>Cribrononion</u> species (=<u>Elphidium</u> species in Bartlett, 1966) require salinities higher than 20 ^O/oo to reproduce. This suggests that these forms will be confined to the lower parts of estuaries where salinities are higher.

Extensive research concerning the arenaceous species <u>Ammotium cassis</u> has been conducted in recent years (Olsson, 1976; Scott <u>et al.</u>, 1977). A study of the Miramichi River estuary suggests that the most abundant <u>A. cassis</u> populations are found where transition zones exist. The "ponding effect" occurring in this zone is caused by altered circulation patterns resulting from river and tidal influences becoming more balanced in the estuary (Scott <u>et</u> <u>al.</u>, 1977). One effect of this is the increase of suspended particulate matter (SPM) which is associated with increased river discharge (Nichols, 1977). This increase of SPM along with rapid sedimentation and increased organic content in the sediments favor the proliferation of <u>A. cassis</u> (Olsson, 1976; Scott et al., 1977).

CHAPTER III

METHODS

Collection of Cores:

Cores for this project were collected during May, 1979, by Medioli and Scott aboard the C.S.S. <u>Navicula</u> operating from Wallace Basin. Approximate positions of Kranck's (1972) Baie Verte core locations were reached and recorded (radar was used for navigation, Refer to Figure 1, Table 1). Other coring locations in the area were sampled but only the original two cores in Kranck's former locations proved useful.

A 600 kilogram Alpine head (with a 3.5 metre barrel) was used as a gravity corer. This was a modification of previous plans to use the complete Alpine piston corer since conditions proved too dangerous aboard the C.S.S. <u>Navicula</u> for piston coring. The two cores (called Navicula #1 and Navicula #2) were stored in plastic core liner until they could be split in the laboratory.

Splitting of Cores For Sampling:

The cores were split according to standard procedures; both the working and the archives halves are stored at Dalhousie University. After splitting, the cores were sampled at 10 cm. intervals (15 c.c. samples). Sometimes closer interval sampling was required, depending on lithology.

Sample Preparation:

All samples were prepared by standard procedures (Scott, 1977; Scott <u>et al</u>., 1977) which will briefly be described here. Each 15 c.c. sample was wet sieved through .5 mm. and .063 mm. sieves with the .5 mm. sieve retaining the coarse organics and allowing foraminifera to pass to the .063 mm. sieve where they were retained. Decantation was used to separate the fine organic material from the microfossils and sediment. The foraminifera samples were preserved in alcohol.

Due to excessively large numbers of foraminifera and volumes of sediment present in most of the samples, only a few of them were examined in the alcohol medium. The remaining samples were dried and the foraminifera separated from the sand by flotation in carbon tetrachloride (sp.g.-1.58). The separated foraminifera were then examined in a dry state. Because of the large numbers of foraminifera found in most of these samples, they were split to a workable number of 300-500 when necessary using an "Otto" microsplitter.

Photography of Specimens:

Those foraminiferal species which were considered significant in the interpretation of the data were photographed using the Cambridge Scanning Electron Microscope located at the Bedford Institute of Oceanography. Polaroid N/P 55 film was used. The photographs were then assembled on a plate and rephotographed for display in this study.

CHAPTER IV

RESULTS

General Observations of Foraminiferal Distributions in the Cores:

Twenty-four species of foraminifera occur in the two cores (Table 2). There are commonly six to twelve of these species in each sample. Many species are represented by only a few individuals. Because of this, just the significant species are considered in the interpretation and correlation (Fig. 2, 3).

Most of the samples in both cores contain very large populations of foraminifera, typically ranging from 2,000 to 5,000 individuals per 15 c.c. Exceptions to this occur in Navicula #2 where two areas show a distinct decline in population size to about 100 individuals. One area occurs just above the bedrock (96 to 86 cm) while the other is located from 42 to 34 cm.

Sedimentary, Foraminiferal, and Environmental Descriptions:

Although a detailed sediment analysis was not conducted, basic distinctions were recorded in the preliminary examination of the cores (Fig. 2, 3).

Navicula #1:

At 15.51 metres below present MSL the base of the unconsolidated sediments is in contact with a bedrock





Lithostratigraphy and foraminiferal occurrences in Navicula Core #1



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Figure 3

Lithostratigraphy and foraminiferal occurrences in Navicula Core #2

of red sandstone (151 cm. in the core, Fig. 2). This is followed by a soil horizon (151 to 142 cm.) which contains no foraminifera and separates the sandstone from the overlying marine deposits. At the base of the marine sequence is an organic layer mixed with wood material (142 to 134 cm.). This peat layer can be divided into two faunal assemblages. The lower assemblage (142-140 c.m) consists of a relatively small population of <u>Protelphidium orbiculare</u> and <u>Cribrononion</u> <u>excavatum</u>. These species are characteristic of an intertidal to subtidal open bay environment. The upper faunal zone (140 to 134 cm.) contains a large arenaceous population dominated by <u>Trochammina inflata</u> with smaller numbers of <u>Tiphotrocha comprimata</u> and <u>Trochammina lobata</u>. This indicates a salt marsh environment.

Above the organic material is a red sand layer (135 to 130 cm.) with an oyster shell horizon extending to the 125 cm. mark. A thick bed of light grey mud (125 to 62 cm.) overlies the shell horizon. From the base of the red sand layer (135 cm.) to the 84 cm. mark in the light-gray mud, an assemblage of <u>P. orbiculare and C. excavatum</u> is found. The upper part of this zone (105 to 84 cm.) contains <u>Ammotium</u> <u>cassis</u> and <u>Eggerella advena</u> which show a gradual increase in numbers approaching the top of the assemblage. Small numbers of <u>Buccella frigida</u> are also present throughout this zone. The predominant calcareous forms suggest an open bay environment which is changing toward the top of the zone.

A sequence of different coloured muds (62 to 0 cm.) overly the light-gray mud and trend from gray on the bottom to red-gray on the top. From the 84 cm. mark in the light-gray mud to the top of the core, as assemblage composed almost entirely of <u>A. cassis</u> and <u>E. advena</u> occurs. These two species (especially <u>A. cassis</u>) indicate a transition zone in an estuary.

Navicula #2:

The basic sediment sequence here is similar to that of Navicula #1, although compressed. At 9.96 metres below present MSL (96 cm. in the core), a bedrock of red sandstone is in contact with an organic layer (Fig. 3). This peat layer (96 to 91 cm.) defines the limits of a faunal assemblage zone. The foraminiferal population is very small here. <u>Trochammina inflata</u> is the most abundant species, with only slightly lesser numbers of <u>Protelphidium orbiculare</u> and <u>Cribrononion excavatum</u>. The strong presence of the <u>T. inflata</u> indicates a salt marsh environment.

Above the peat is a relatively thick gray sand and oyster shell horizon (91 to 65 cm.). Light gray mud extends from the top of the shell layer to the 26 cm. mark. From the bottom of the oyster shell layer to the middle of the light gray mud layer (91 to 40 cm.) a calcareous fauna is dominant. <u>P. orbiculare and C. excavatum are both very prolific while Buccella frigida and Cribrononion bartletti occur in minor</u> but persistent numbers. One small section from 64 to

62 cm. contains a significant population of <u>C. gunteri</u>. The abundance of calcareous forms suggests that this is an open bay environment.

Overlying the light gray mud, a series of coloured muds occur (65 to 0 cm.) with tan to light-gray on the bottom grading into red-gray at the top of the core. An assemblage dominated by <u>Ammotium cassis</u> and <u>Eggerella advena</u> occurs at the 40 cm. mark in the light-gray mud and continues to 0 cm. A transition zone is indicated by the presence of this assemblage.

CHAPTER V

DISCUSSION

General Comments:

The peat layers in the Navicula cores are presently being C-14 dated; however, the results are not available at the present time to be submitted with this study. McRoberts (1968) and Kranck (1972) publisheā C-14 dates which should correspond quite closely to the Navicula cores (Table 1), with one exception. The 10 metre core in the present study should be slightly younger than McRoberts' 11.8 metre core. Based on a proportionality calculation of sea level rise using McRoberts' (1968) C-14 date, it can be inferred that the peat layer at 10 metres below present MSL (in the present study) occurred approximately 2,500 years B.P. With these assumptions in mind, the dates discussed in this study will be approximations based on the results of McRoberts (1968) and Kranck (1972).

Faunal Assemblage Comparisons With Modern Day Environments:

Both cores show the same vertical sequence of faunal assemblages with one exception. Navicula #1 has a small open bay zone at the base which is not found in Navicula #2. Other than this, both cores have a marsh foraminiferal assemblage at the bottom overlain by an open shallow bay

assemblage and an estuarine transition zone occuring at the top.

The zone identified as the marsh environment contains a fauna similar to that described by Scott and Medioli (1978a, 1980). The important species are <u>Trochammina inflata</u> and <u>Tiphotrocha comprimata</u>. These arenaceous forms are dominants found in the upper marsh zones of the Maritimes (Scott and Medioli, 1978a, 1980).

This zone (in both cores) shows a relatively dramatic decrease in individuals at the base, which is strongly characteristic of higher high water areas in a marsh (Scott and Medioli, 1980).

<u>Cribrononion excavatum</u> and <u>Protelphidium orbiculare</u> define the open bay assemblage zone. These calcareous species are strongly characteristic of an intertidal to shallow, subtidal environment (Scott <u>et al</u>., in press). This assemblage was observed by Scott <u>et al</u>. (1977) in the Miramichi River estuary. A salinity of greater than 20 o/oo appears to be the requirement for the proliferation of these species (Bartlett, 1966). These conditions are satisfied in the open bay zone of an estuary.

Large populations of Eggerella advena and Ammotium cassis are strongly characteristic of an estuarine transition zone. This assemblage has been identified by Scott et al., (1977) in the Miramichi River estuary. With the aid of cluster analysis, they suggested that only <u>A. cassis</u> can

define this zone while <u>E.</u> advena can be found in large numbers in either the transition or open bay faunal assemblage.

Paleoecology:

The evidence presented distinctly implied that the Baie Verte estuary developed from a two assemblage zone to a three zone system. It should be noted that the upper estuarine assemblage zone (defined by <u>Miliammina fusca</u> and and <u>Ammotium salsum</u>, Scott <u>et al.</u>, 1977) described by Scott <u>et al.</u>, (1977) was not found in the present study. This probably indicates that it was poorly developed due to low fresh water input, similar to conditions exhibited in Prince Edward Island estuaries (Scott et al., in press).

In the early phase of the estuary, it consisted of a fringing marsh zone and a directly adjacent shallow open bay An arenaceous fauna of Trochammina inflata and zone. Tiphotrocha comprimata dominated the marsh assemblage while open bay conditions were represented by the calcareous species Protelphidium orbiculare and Cribrononion excavatum. The absense of a transition zone was probably due to a circulation imbalance, with the marine influences being far greater than river influences (Scott et al., 1977). With time there was a change from this nontransitional estuary to a transitional type. This was apparently caused by the " ponding effect" which most likely resulted from more balanced marine and fluvial influences due to increased river discharge and/or decreased tidal influence-from rising

sea level (Scott <u>et al.</u>, 1977). The result was a transition assemblage zone characterized by <u>Ammotium cassis</u> and <u>Eggerella</u> <u>advena</u> separating the fringing marsh from the more seaward open bay environment. It should be noted here that the assumption has been made that an open bay zone presently exists in a seaward direction from the study area cores.

Implications Regarding Relative Sea Level Changes:

The foraminiferal distributions indicate that 5,000 years B.P. (approximation of C-14 date, Kranck, 1972), there was a short emergence stage in the Baie Verte estuary at 15.5 metres below present MSL. This is indicated by the open bay assemblage at the base of Navicula #1, followed by a marsh sequence (suggesting falling sea level). A marine transgression then began. At approximately 2,500 years B.P. (calculated from C-14 date, McRoberts, 1968) the marine water had reached a point 10 metres below present MSL. The evidence indicates that this submergence has apparently been continuous to the present day.

Relationship Between Relative Sea Level Changes and Deglaciation:

The results appear to conform with the studies of the migration of the "peripheral bulge" in the Maritimes (e.g. Kranck, 1972; Scott, 1977). The temporary regression of the sea at the base of Navicula #1 would suggest that the maximum point of emergence is this area (where the cores were obtained) was approximately 5,000 years B.P. at 15.5

metres below present MSL. Since that time there has been continuous submergence. These results would be consistent with the suggestion that the "peripheral bulge" has already migrated westward through the western part of the Northumberland Strait and into New Brunswick (Scott, 1977). Thus, the last effect of the "bulge" in the Baie Verte area occurred approximately 5,000 years B.P. with the emergent feature of the salt marsh being followed by submergence.

Paleoclimatic Conditions:

Most of the species (including Protelphidium orbiculare, Cribroelphidum excavatum, Eggerella advena, etc.) in the cores are associated with temperate to cold temperate waters (Bartlett, 1966). This suggests that the climatic conditions over the last 5,000 years have probably been quite similar to the cool, temperate conditions which exist in the Northumberland Strait today. One exception to this assumption is the appearance of Cribrononion gunteri in Navicula #2 (64 to 62 cm.). This species is associated with warm temperate to subtropical waters (Snyder and Katrosh, 1979). It might be suggested that conditions in the Northumberland Strait were warmer at one time during the last 5,000 years. However, because of the complexity of this body of water and insufficient studies done in the area, any conclusions drawn must be considered as mere speculation.

Comparison to McRobert's (1968) Work:

A comparison of this study with that done by McRoberts (1968) indicates a great deal of variation in results obtained in the same study area. Although the lithologies recorded in the two studies are very similar, the foraminiferal distributions show wide discrepancies. To begin with, her results show the basal peat layer as depauperate of foraminifera while the Navicula cores have significant numbers of arenaceous foraminifera in the equivalent layers. The implication here is that the peat was not a fresh water bog, but definitely the remains of a salt marsh formation. Thus, the organic material can be very accurately used as an indicator of former sea level (Scott, 1977; Scott and Medioli, 1978a, 1980).

Another point of difference is that McRoberts (1968) described the upper section of the core as either barren or depauperate. In the Navicula cores very large populations of arenaceous species are present (to the top of the cores). This would suggest that the foraminiferal fauna is flourishing in this area at the present day and not depauperate as suggested by McRoberts (1968).

The differences encountered in the two studies can probably be accounted for in terms of sampling techniques. The methods for preserving the foraminifera and extracting them from the cores have been substantially refined in very recent years. Because many of these techniques were

not available at the time of the McRoberts (1968) study, it can be suggested that possibly the large arenaceous populations (observed in the present study) were destroyed when the cores were processed or sampled.

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CHAPTER VI

CONCLUSIONS

(1) The organic layer at the base of each core contains an abundant arenaceous population including <u>Trochammina inflata</u> and <u>Tiphotrocha comprimata</u>. This confirms that these peat layers were salt marsh deposits and can be used as accurate indicators of first invasions of a marine environment.

(2) Foraminiferal distributions found in the cores suggest that the initial Baie Verte estuary in the study area consisted of two faunal assemblage zones. <u>Trochammina inflata</u> and <u>Tiphotrocha comprimata</u> define the fringing marsh environment while the calcareous forms <u>Protelphidium orbiculare</u> and Cribrononion spp. indicate the open bay environment.

(3) The most recent sediments in the study area indicate the presence of a transition zone in the estuary which is identified by large populations of <u>Ammotium cassis</u>.

(4) The results suggest that there has been a shift in the Baie Verte estuary from a nontransitional, two-zoned system to the present conditions of a transitional, three-zoned estuary.

(5) Relative sea level changes indicate that approximately 5,000 years B.P., there was a transgression of the sea in the Baie Verte estuary. This was followed by a period of emergence and finally a continuous submergence to the present day.

(6) This study provides evidence supporting the recent models of the migration of the "peripheral bulge" in the Maritimes. The results suggest that the "bulge" has already passed through the western part of the Northumberland Strait so that the Baie Verte area is now located at the distal end. The maximum emergence (the effect of the "bulge") in this area was experienced approximately 5,000 years B.P. at 15.5 metres below present MSL.

(7) The species of foraminifera identified in the cores indicate that the paleoclimatic conditions over the last 5,000 years have most probably been similar to the cool, temperate conditions which exist at present.

(8) A comparison with results obtained by McRoberts (1968) indicates variations from the data presented in this study concerning the same area. The basic difference is that McRoberts found the peat layers and large sections of the top of the cores barren of foraminifera while in the present study, both of these sections contained abundant arenaceous species. there a

CHAPTER VII

SYSTEMATIC TAXONOMY

Of the foraminifera identified in this study, only those species which showed significant occurrences will be presented here. The references cited consist of original references and those that were pertinent to this study. The generic names are in accordance with Loeblich & Tappan (1964).

Ammonia beccarii (Linné)

<u>Nautilus beccarii</u> Linné. 1758, p. 710.
<u>Ammonia beccarii</u> (Linné). Brunnich, 1772, p. 232; Frizzell
and Keen, 1949, p. 106; Gregory, 1970, p. 222, Pl. 12, figs.
4-6; Schnikter, 1974, pp. 216-223, Pl. 1; Cole and Ferguson,
1975, p. 32, Pl. 9, figs. 1, 2; Scott, 1977, pp. 164-165,
Pl. 6, figs. 10, 11; Schafer and Cole, 1978, p. 27, Pl. 8,
fig. 6; Scott and Medioli, 1980, p. 35, Pl. 5, figs. 8, 9.
<u>Streblus beccarii</u> (Linné). Fischer de Waldhiem, 1917, p. 449,
Pl. 13; Bradshaw, 1957, p. 1138, text fig. la-c.
<u>"Rotalia" beccarii</u> (Linné) var. <u>tepida</u> Cushman; 1926, p. 79,
Pl. 1; Parker, 1952a, p. 457, Pl. 5, figs. 7, 5.

Ammotium cassis (Parker)

Plate I, figure 7

Lituola cassis Parker in Dawson (1870) pp. 177, 180, fig. 3.

Buccella frigida (Cushman). Anderson, 1952, p. 144, figs. 4 a-c, 5, 6 a-c; Gregory, 1970, p. 220, Pl. 12, figs. 1-3; Cole and Ferguson, 1975, p. 33, Pl. 8, figs. 8, 9; Scott, 1977, p. 167, Pl. 8, figs. 10, 11; Scafer and Cole, 1978, p. 27, Pl. 8, figs. 1, 2.

Cribrononion bartletti (Cushman)

Plate I, figure 16

Elphidium bartletti.Cushman, 1933, p. 4, Pl. 1, fig. 9; Gregory, 1970, p. 225, Pl. 13, figs. 3-5; Cole and Ferguson, 1975, p. 34, Pl. 7, figs. 3, 4; Schafer and Cole, 1978, p. 27, Pl. 10, fig. 4.

Cribroelphidium bartletti (Cushman). Scott, 1977, p. 169, Pl. 6, fig. 8.

Scott and Medioli (1980) have suggested that generic distinctions between Elphidium, Cribroelphidium, and Cribrononion require clarification for their proper usage. First of all, they do not agree with the criterion established by Loeblich and Tappan (1964) as acceptable in differentiating between Cribroelphidium and Cribrononion. The distinction that Loeblich and Tappan (1964) made was that Cribroelphidium has areal apertures in addition to the row of pores at the base of the septal face while Cribrononion lacks these areal apertures. According to Scott and Medioli (1980), this does not provide an acceptable basis for making a generic distinction; therefore, if the two can be considered as synonyms, the genus name Cribrononin has priority and must be retained. In the particular species encountered in this study, <u>Elphidium</u> does not appear appropriate either, according to the narrow definition invoked by Loeblich and Tappan (1964). The result is that <u>Cribrononion</u> will be the generic name used where deemed necessary (Scott and Medioli, 1980).

Cribrononion excavatum (Terquem)

Plate I, figures 14, 15

Polystomella excavata Terquem, 1876, p. 429,Pl. 2, figs. 2a-d.
Polystomella striato-punctata (Fichtel and Moll) var.
selseyensis.Heron - Allen and Earland, 1911, p. 448.
Elphidium excavatum (Terquem). Cushman, 1930, p. 21, Pl. 8,
figs. 1-3; Schafer and Cole, 1978, p. 27, Pl. 9, fig. 7;
Levy <u>et al</u>., 1975, p. 176, fig. 9, Pl. 3, figs. 1, 2, 5, 6.
Cribrononion excavatum (Terquem). Lutze, 1965, pp. 96-101, Pl.
15, fig 39; Lévy <u>et al</u>. 1969, p. 93, Pl. 1, figs.
1a, b, 2a, b, 4a, b; Scott and Medioli, 1980, p. 35, Pl. 5,
figs. 5, 6.

Elphidium excavatum (Terquem) formae. Feyling - Hanssen, 1972, pp. 337-354, Pls. 1-6.

<u>Cribroelphidium excavatum</u> (Terquem) forma <u>clavatum</u> (Cushman). Scott <u>et al.</u>, 1977, pp. 1578, 1579, Pl. 5, figs. 1, 2; Scott, 1977, p. 169, 170, Pl. 6, fig. 2.

Cribroelphidium excavatum (Terquem) forma selseyensis (Heron-Allen and Earland). Scott <u>et al</u>., 1977, p. 1579, Pl. 5, fig. 3; Scott, 1977, p. 170, Pl. 6, fig. 3.

Both formae, clavatum and selseyensis, are considered

under this species because it has been suggested that any distinction between the two in the past was arbitrary (Scott and Medioli, 1980). This decision has been substantial as a result of photographic documentation by Miller (1979) showing an intergradational series between the two formae.

Cribrononion gunteri Cole

Plate I, figure 13

Elphidium gunteri.Cole, 1931, p. 34, Pl. 4, figs. 9, 10; Phleger, 1960, Pl. 3, fig. 22; Snyder and Katrosh, 1979, Pl. 2, figs. 1, 2

Elphidium gunteri Cole, var. galvestonense Kornfeld; Phleger and Parker, 1948, p. 10, Pl. 5, fig. 14 Ephidium gunteri Cole, forma typicum. Poag, 1978, p. 402, Pl. 2, figs. 13-16.

Investigations by Poag (1978) have resulted in <u>C</u>. <u>gunteri</u> (<u>E gunteri</u> according to Poag) being classified as a distinct ecophenotype. However, more recent work done by Miller and Scott indicates that it would be more appropriate to consider <u>C</u>. <u>gunteri</u> as an ecophenotype of <u>C</u>. excavatum (Miller & Scott, personal communication, 1980).

Cribrononion subarcticum (Cushman)

Plate I, figure 12

Elphidium subarcticum.Cushman, 1944, p.27, Pl. 3, figs.34-35; Parker, 1952a, p. 449, Pl. 4, figs. 3-6, 8; Phleger and Walton, 1950, p. 277, Pl. 2, figs. 19, 20; Gregory, 1970, p. 229, Pl. 14, fig. 7; Cole and Ferguson, 1975, p. 34, Pl. 8, figs. 1, 2; Schafer and Cole, 1978, p. 27, Pl. 10, fig. 1. <u>Cribroelphidium subarcticum</u> (Cushman). Scott, 1977, p. 171, Pl. 6, fig. 7.

Eggerella advena (Cushman)

Plate I, figure 8

Verneuilina advena.Cushman, 1921, p. 141.

Eggerella advena (Cushman). Cushman, 1937, p. 51, Pl. 5, figs 12-15; Phleger and Walton, 1950, p. 277, Pl. 1, figs. 16-18; Parker, 1952b, p. 404, Pl. 3, figs. 12, 13; Parker, 1952a, p. 447, Pl. 2, fig. 3; Gregory, 1970, p. 183, Pl. 4, figs. 1-3; Cole and Ferguson, 1975, p. 34, Pl. 3, figs. 10, 11; Scott <u>et al</u>., 1977, p. 1579, Pl. 2, fig. 7; Scott, 1977, p. 171, Pl. 6, fig. 9; Schafer and Cole, 1978, p. 27, Pl. 3, fig. 1; Scott and Medioli, 1980, p. 40, Pl. 2, figure 7.

Protelphidium orbiculare (Brady)

Plate I, figure 17

Nonionia orbiculare.Brady, 1881, p. 415, Pl. 21, fig. 5.
Nonion orbiculare (Brady). Cushman, 1930, p. 12, Pl. 5, figs.
1-3.

Elphidium orbiculare (Brady). Hessland, 1943, p. 262; Gregory, 1970, p. 228, Pl. 14, figs. 5, 6.

<u>Protelphidium orbiculare</u> (Brady). Todd and Low, 1961, p. 20, Pl. 2, fig. 11; Cole and Ferguson, 1975, p. 39, Pl. 7, figs. 7, 8; Scott et al., 1977, p. 1579, Pl. 5, figs. 5,6; Scott 1977, pp. 174, 175, Pl.6, fig. 9; Shafer and Cole, 1978, p.28, Pl. 10, fig. 5; Scott and Medioli 1980, p. 43, Pl. 5, fig. 7.

Reophax arctica Brady

<u>Reophax arctica</u>.Brady, 1881, p. 405, Pl. 21, fig. 2 a, b;
Parker, 1952b, p. 395, Pl. 1, figs. 6, 7; Gregory, 1970,
p. 168; Pl. 2, fig. 3; Cole and Ferguson, 1975, p. 40,
Pl. 1, fig. 9; Scott, 1977, p. 175, Pl. 3, fig. 5; Schafer
and Cole, 1978, p. 29, Pl. 2, fig. 5.

Bigenerina artica (Brady). Cushman, 1948, p. 31, Pl. 3, fig. 9.

Reophax nana Rhumbler

<u>Reophax nana</u>. Rhumbler, 1911, p. 182, Pl. 8, figs. 6-12;
Parker, 1952a, p. 457, Pl. 1, figs. 14, 15; Scott <u>et al.</u>,
1977, p. 1579, Pl. 3, figs. 1, 2; Scott, 1977, p. 175,
Pl. 3, fig. 7; Schafer and Cole, 1978, p. 29, Pl. 2, fig. 4;
Scott and Medioli, 1980, p. 43, Pl. 2, fig. 6.

Reophax nodulosa, Brady

Plate I, figure 9

<u>Reophax nodulosa</u>, Brady, 1879, p. 52, Pl. 4, figs. 7, 8; Brady, 1884, p. 294, Pl. 31, figs. 1-9; Schafer and Cole, 1978, p. 29, Pl. 2, fig. 6.

Reophax scorpiurus, de Montfort

<u>Reophax scorpiurus</u> de Montfort 1808, p. 330; Scott <u>et al.</u>, 1977, p. 1579, Pl. 3, figs. 3, 4.

Tiphotrocha comprimata (Cushman and Bronnimann)

Plate I, figures 3, 4

Trochammina comprimata. Cushman and Brönnimann, 1948b, p. 41, Pl. 8, figs. 1-3.

<u>Tiphotrocha comprimata</u> (Cushman and Brönnimann). Saunders, 1957, p. 11; Parker and Athearn, 1959, p. 341, Pl. 50, figs. 14-17; Scott <u>et al.</u>, 1977, p. 1579, Pl. 4, figs. 3,4; Scott, 1977, p. 176, Pl. 5, figs. 14-16; Scott and Medioli, 1980, p. 44, Pl. 5, figs. 1-3.

Trochammina inflata (Montagu)

Plate I, figures 1,2

<u>Nautilus inflatus</u>. Montagu, 1808, p. 81, Pl. 18, fig. 3.
<u>Trochammina inflata</u> (Montagu). Parker and Jones, 1859,
p. 347; Phleger and Walton, 1950, p. 280, Pl. 2, figs. 1-3;
Parker, 1952b, p. 407, Pl. 4, figs. 6, 10; Parker, 1952a,
p. 459, Pl. 3, fig. 2a, b; Gregory, 1970, p. 180, Pl. 4,
figs. 3, 4; Cole and Ferguson, 1975, p. 43, Pl. 4, figs.
3, 4; Scott, 1977, p. 177, Pl. 4, figs. 12-14, Pl. 5, figs.
1-3; Scafer and Cole, 1978, p. 29, Pl. 5, fig. 2; Scott and
Medioli, 1980, p. 44, Pl. 3, figs. 12-14, Pl. 4, figs. 1-3.

Trochammina lobata Cushman

Trochammina lobata.Cushman, 1944, p. 18, Pl. 2, fig. 10; Phleger and Walton, 1950, p. 281, Pl. 2, figs. 4, 5; Parker 1952b, p. 408, Pl. 4, fig. 7a, b; Parker, 1952a, p. 459, Pl. 3, fig. 2a, b; Gregory, 1970, p. 180, Pl. 4, figs. 5, 6; Cole and Ferguson, 1975, p. 43, Pl. 4, figs. 5, 6; Scott <u>et</u> <u>al.</u>, 1977, p. 1579, Pl. 4, figs. 1, 2; Scott, 1977, p. 179, Pl. 5, figs. 8, 9; Schafer and Cole, 1978, p. 29, Pl. 4, fig. 5.

Trochammina macrescens Brady

Plate I, figures 5, 6

Trochammina inflata (Montagu) var. macrescens.Brady, 1870, p. 290, Pl. 11, fig. 5a-c; Scott, 1976, p. 320, Pl. 1, figs. 4-7; Scott <u>et al.</u>, 1977, p. 1579, Pl. 4, figs. 6, 7; Scott, 1977, p. 178, Pl.178, Pl. 4, figs. 1-8.

<u>Jadammina polystoma</u>. Bartenstein and Brand, 1938, p. 381, figs la-c, 2a-l; Parker and Athearn, 1959, p. 341, Pl. 50, figs. 21, 22, 27; Scott, 1977, p. 173, Pl. 4, figs. 9-ll. <u>Trochammina macrescens</u> Brady. Phleger and Walton, 1950, p. 281, Pl. 2, figs. 6, 7; Parker, 1952b, p. 408, Pl. 4, figs. 8a, b; Parker, 1952 a, p. 460, Pl. 3, figs. 3a, b; Parker and Athearn, 1959, p. 341, Pl. 50, figs. 23-25; Gregory, 1970, p. 181, Pl. 4, fig. 7; Cole and Ferguson, 1975, p. 43, Pl. 4, figs. 6, 7; Schafer and Cole, 1978, p. 29, Pl. 4, fig. 3; Scott and Medioli, 1980, p. 44, Pl. 3, figs. 1-8.

Jadammina macrescens (Brady). Murray, 1971, p. 41, Pl. 13, figs. 1-5.

It should be noted that the poor condition of the

individuals, as seen in the accompanying photographs, was typical of this species in the two cores examined.

<u>Trochammina squamata</u> Parker and Jones <u>Trochammina squamata</u>.Parker and Jones, 1865, p. 407, Pl. 15, figs. 30, 31a-c; Phleger and Walton, 1950, p. 281, Pl. 2, figs. 12, 13; Parker, 1952b, p. 408, Pl. 4, figs. 11-16; Parker, 1952a, p. 460, Pl. 3, fig. 4a, b; Cole and Ferguson, 1975, p. 43, Pl. 4, figs. 11, 12; Scott, 1977, p. 180, Pl. 5, figs. 6, 7; Schafer and Cole, 1978, p. 29, Pl. 5, fig. 1; Scott and Medioli, 1980, p. 45, Pl. 4, figs. 6, 7.

Plate I

Figure	1,2.	Trochammina inflata (Montagu) l. dorsal view, $76.4 \times 2.$ ventral view, $76.4 \times .$
Figure	3,4.	Tiphotrocha comprimata (Cushman and Brönnimann) 3. dorsal view, 100.9 x, 4. ventral view, 93.5 x.
Figure	5,6.	$\frac{\text{Trochammina}}{69.4 \text{ x, 6. ventral view, }} \text{ Brady 5. dorsal view, }$
Figure	7.	Ammotium cassis (Parker) 7. side view, 53.5 x.
Figure	8.	Eggerella advena (Cushman) 8. side view, 83.7 x.
Figure	9.	Reophax nodulosa Brady 9. side view, 62.5 x.
Figure	10,11.	Buccella frigida (Cushman) 10. dorsal view, 53.9 x , 11. ventral view, 75.2 x.
Figure	12.	Cribrononion subarcticum (Cushman) 12. side view, 102 x.
Figure	13.	Cribrononion gunteri Cole 13. side view, 72.7 x.
Figure	14,15.	Cribrononion excavatum (Terquem) 14. side view, $65.4 \times 15.$ side view, 50.7×10.7
Figure	16.	Cribrononion bartletti (Cushman) 16. side view. 37.2 x.
Figure	17.	Protelphidium orbiculare (Brady) 17. side view, 60.5 x.



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APPENDIX

Τ.	AB	LE	Ξ	1

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Description of Comparison of Cores

Navicula Core #	Latitude and Longitude	Water Depth (m)	Core Length to base of organic layer (m)	Corresponding McRoberts (1968) Core #	* Corresponding** Kranck (1972) Core #	Water Depth (m)	Core Length (m)	Carbon 14 Dates for McRoberts (1968) and Kranck (1972) (years B.P.)
1	45 ⁰ 57.8, N 63 ⁰ 42.8 W	14.0	1.4		4	14.0	1.2	5125 [±] 120 (Kranck, 1972)
2	46 ⁰ 2.2. N 63 ⁰ 48 W	9.0	1.0	11	2	9.0	2.8	3000 ± (McRoberts, 1963)

- * McRobert's (1968) core #11 and Kranck's (1972) core #2 are the same; therefore, only one description is necessary for both.
- ** The Navicula cores are the same as McRoberts' (1968) core and Kranck's (1972) cores according to water depth rather than map coordinates.

TABLE 2

																		-												
CORE NUMBER AND DEPTH (cm.)	1-0	1 -14	1-22	1-34	1-44	1-54	1-64	1-76	1-84	1-94	1-104	1-118	1128	1-134	1-137	1-140	2-0	2-10	2-20	2-28	2-34	2-40	2-50	2-58	2-62	2-70	2-75	2-80	2-85	2-91
No. of species	6	6	6	6	9	10	6	3	13	9	12	8	8	12	11	7	7	12	5	5	7	7	9	7	7	7	S	7	7	8
No. of Individuals per 15 cc	4032	2040	2402	5312	5674	754	7560	3168	1708	2216	2000	2056	7230	430	3201	437	2812	4232	8768	1460	103	100	640	1832	2840	2380	4880	384	106	60
Ammobaculites dilatatus															x															
Arronia becarii																														2
Amentium cassis	23	4	50	36	51	66	58	58	40	27	22	1		1			27	22	28	13	23	30	6							
A. salsum						-					X				1															
Puccella frigida						1			2	8	1	5	6	X	X	4							4	8	5	10	12	8	10	2
Cribrononion bartletti						3			1	1	1	2	9			7							3	4	6	2	2	5	11	;
C. excavatum					X	13			26	27	25	33	36	X	4	26		X			1	1	26	36	34	36	49	53	25	18
C. gunteri			1	1																					2			1	1	
C. subarcticum		1									X		4										X		1	i	6	2	9	
Eggerella advena	7ó	93	49	61	44	3	39	38	19	10	18	1		X		X	67	75	69	84	50	63	5							2
Hoplostragnoides bonolanti														х																
Perisphaenammina brady																		Х												
Proteinulaium orbitulare					2	13	1		10	24	25	57	4.3	х	3	59						۱	52	50	52	49	30	29	36	26
Fseudorwiy Kurphina rovergijae						x							X																[
Reopraz arctica	X	X		1	Х				X																					
R. nana		1		X	1		X		X		X							X		X	+	1								
R. nodulosa	X	Х	X		1	Х	1		X	1								Х	X										1	
P. scorolu rus		1			Х	2			. X								1	X												
Spiroplectammina biformis	X		X											×	x		х	x												
Tiphotrocha comprimata				1		X			X		1	x		24	26			x			4			x			х	1	5	2
Trocharmina inflata			X							x	1		١	52	63	4	x	x		X	5		2	1		1	x		2	34
T. lobata		1	1											12	1		1											2		
T. macrescens									X			X		7	1			3			1	1						1		13
T. squamata	1	2	X	2	X		1	4	1	2	5		X	1	X	X	3	1	2	z	15	3	1	1	X	X	X			

Table 2. For aminiferal percentages in Navicula Cores 1 and 2; X is less than 1% of total

The following is a breakdown of the time spent in preparation of this thesis.

Item	Time Spent	(Hrs)
Foraminifera Identification	120	
Sample Preparation	10	
Tables and Diagrams	15	
Touching up of Photographs	4	
Preparation of Photographic Plate	l	
Research and Writing	200	
Total Time	350	

Dave Scott and Bhan Deonairine did the SEM photography at BIO.

The final draft of two diagrams and one table was done by the Geology Department.

The typist did the final draft of the other table.

All of coring and most of sample preparation was done by Drs. Scott and Medioli.

Garth Prime