

Analytical programme and applied methods

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The laboratory phase of the 'Nordolie' project included geochemical, microscopical and petrophysical methods (fig. 9). These methods vary considerably in approach and degree of sophistication. Laboratory work has been carried out in Copenhagen at the laboratories of the Geological Survey of Greenland (GGU), the Geological Survey of Denmark (DGU), and the Geological Institute, University of Copenhagen.

Sampling

Considerable time was devoted to stratigraphic, sedimentological and structural studies of potential reservoir and source rock sequences with the highest priority during field work given to sampling of fine-grained units for analytical work. During the summers of 1984 and 1985 two teams carried out field work and sampling from 33 camp sites in central and western North Green-

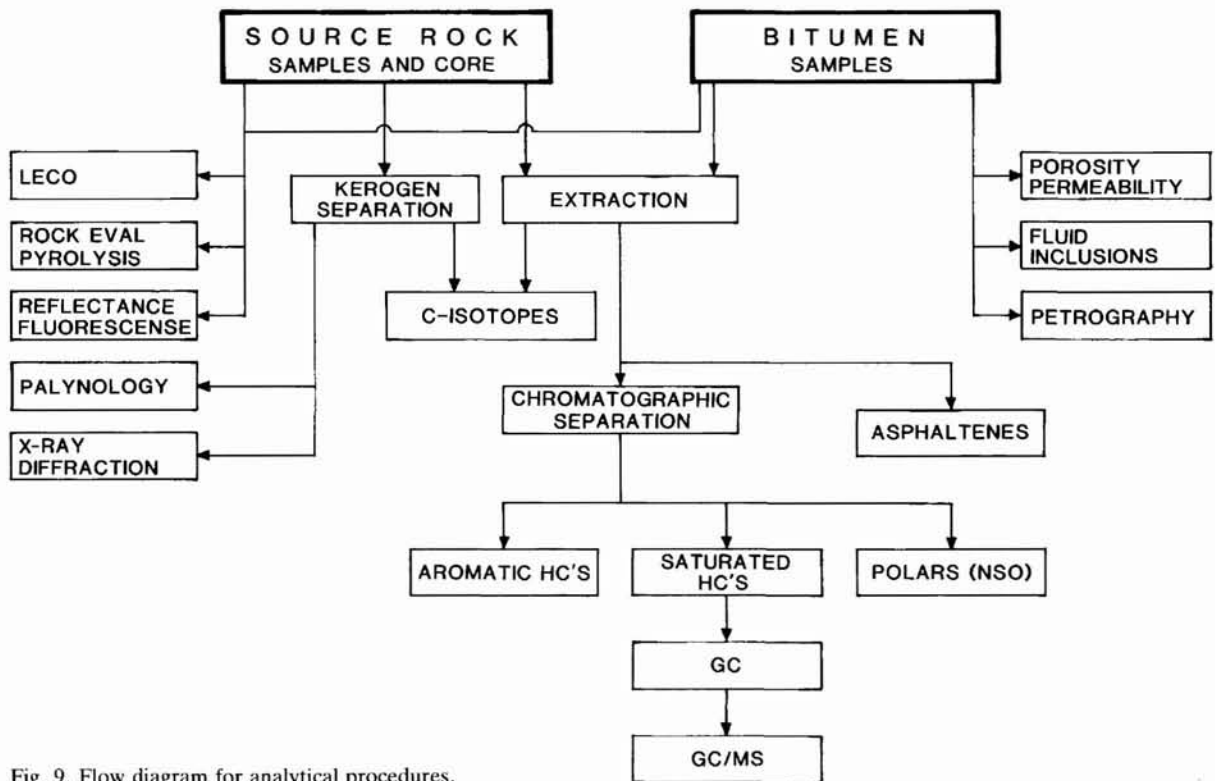


Fig. 9. Flow diagram for analytical procedures.

land (Christiansen & Rolle, 1985; Christiansen *et al.*, 1986); approximately 70 localities were visited by helicopter. In addition to the 989 samples collected by geologists directly involved in the source rock study, a limited number of samples (approximately 75) collected by other geologists were also studied. The study of the Peary Land region, which is only briefly treated in the present work, is mainly based on 275 samples collected by Flemming Rolle in 1980 (Rolle, 1981) plus approximately 50 samples collected by a number of other geologists in 1978 to 1980.

The samples collected were selected to be representative of the studied interval. Whenever possible, samples were collected from unweathered intervals below the surface, as weathering is known to influence the organic geochemistry of surface samples (Leythaeuser, 1973; Clayton & Swetland, 1978; Forsberg & Bjorøy, 1983). All samples were wrapped in aluminium foil to avoid contamination during transport and storage. So far no contaminants have been detected in any of the analysed surface samples.

Drilling programme

A drilling programme was carried out in order to obtain unweathered and statistically representative samples of the most interesting source rock units (see Christiansen *et al.*, 1986 for details). The drilling unit, which was constructed and later modified a number of times by J. Boserup and A. Clausen at GGÜ, has been successfully employed in Jameson Land, East Greenland (Surlyk, 1983; Surlyk *et al.*, 1984), in North Greenland (Christiansen *et al.*, 1986), and on Traill Ø, East Greenland (Marcussen *et al.*, 1987).

Thirteen holes were drilled to a maximum depth of 40 m. Approximately 345 cumulative metres were drilled and except for the uppermost metres at each site the recovery was close to 100%. All cores were described at the drill site using a scale of 1:50 or 1:100 and most holes were logged by gamma ray measurements (logs of the Silurian cores are shown by Christiansen & Nøhr-Hansen, 1989).

After logging, the core pieces were wrapped in aluminium foil and packed in standard core boxes made of hard plastic. Unfortunately, the plastic seems to have caused minor contamination with C₁₂, C₁₄, C₁₆, C₁₈, C₂₀ *n*-alkanes in some of the analysed core samples (see fig. 11).

Handling and storage

All samples and core boxes were transported to Copenhagen by air during or shortly after the two field

seasons. The sampled material is stored at GGU, mostly in closed standard sample cases. During preparation (e.g. cutting, crushing) contact with equipment or containers made of organic compounds (e.g. rubber, plastic) was avoided.

LECO and Rock Eval pyrolysis

The LECO and Rock Eval analyses were carried out at the source rock laboratories at DGU and GGU. More than 600 samples were analysed, mainly during the months immediately following the two field seasons with later supplementary analyses of a smaller number of specific samples. Both types of analyses are based on crushed whole rock material using 200 mg and 100 mg of sample, respectively.

The total carbon content (TC), the total organic carbon content (TOC), and hence the total inorganic carbon content (TIC = TC - TOC) were determined by combustion in a LECO IR 212 furnace before and after treatment with hot concentrated HCl, respectively.

The Rock Eval pyrolysis method (Espitalié *et al.*, 1977) is universally applied by oil and service companies in the characterization of source rock potential, types of organic matter, and thermal maturity (Horsfield *et al.*, 1983; Horsfield, 1984; Tissot & Welte, 1984). However, the method should be used with caution because of variation due to differences in material type and mineral matrix (Espitalié *et al.*, 1980, 1984; Evans & Felbeck, 1983; Katz, 1983) and presence of migrated bitumen (Clementz, 1979). In order to reduce errors and misinterpretations the material from North Greenland was analysed employing the same laboratory procedure and the same instrumentation throughout.

During the Rock Eval pyrolysis the powdered samples are heated at steadily increasing temperatures from 300°C to 550°C. The release curve of the pyrolysate typically shows two peaks (fig. 10). The low temperature peak, S1 (mg HC/g rock) at 300°C is due to hydrocarbons already present in the sample. The second, S2 (mg HC/g rock) between 410°C and 520°C, correlates with hydrocarbons generated by thermal alteration of the kerogen. The CO₂ generated during pyrolysis in the temperature interval 300°C to 390°C is trapped and analysed by thermal conductivity detection as a third peak S3 (mg CO₂/g rock). With increasing thermal maturity the area of the S2 peak decreases and the peak is displaced towards higher temperatures. The temperature of maximum generation (T_{max}) during pyrolysis is used as a maturity parameter (fig. 10).

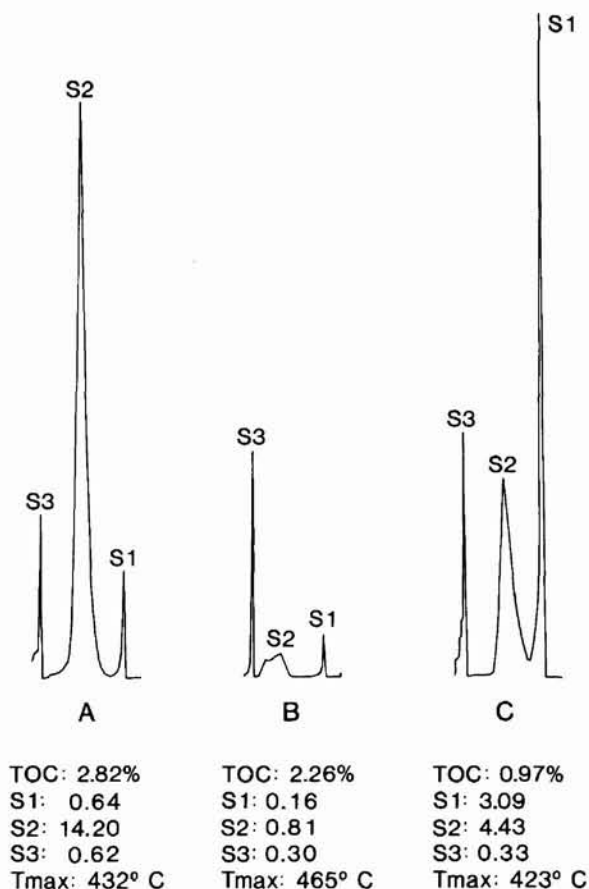


Fig. 10. Selected pyrogrammes showing typical relations of S1, S2, S3, T_{max} and TOC for (A) immature to mature source rock (sample 316775, Lafayette Bugt Formation, Washington Land), (B) mature to postmature source rock (sample 324490, Thors Fjord Member, Warming Land), (C) bitumen-impregnated sandstone (sample 322205, Buen Formation, Wulff Land).

Palynological studies

Palynological preparation and studies were carried out at GGU. Kerogen from more than 455 samples was examined optically in order to obtain information on the Thermal Alteration Index (TAI), kerogen composition, relative kerogen content and content of palynomorphs.

The kerogen was separated from 20 g of each sample by standard palynological preparation, in which minerals are dissolved by HCl and HF. The first slide was made after this acid treatment. The second slide was made of organic residue which had been sieved on a 10 micron nylon mesh. Occasionally a third slide was made

after brief oxidation (3 to 5 minutes) with fuming HNO_3 followed by washing with a weak KOH solution.

The organic residue was mounted in a permanent medium Eukitt® (produced by O. Kindler, West Germany). The first and second slides were used in the evaluation of maturity and kerogen content, the third for biostratigraphic purposes. The slides were studied with translucent microscopy applying an Olympus BH-2 microscope. The S.E.M. observations were carried out on Au-coated sieved and oxidized kerogen using a Cambridge or a Phillips Scanning Electron Microscope at the Geological Institute, University of Copenhagen.

Gas chromatography and gas chromatography/mass spectrometry

Gas chromatography (GC) and gas chromatography/mass spectrometry (GC/MS) analyses were carried out at the source rock laboratories at DGU and GGU. Following preliminary interpretation based on screening methods (LECO/Rock Eval and palynological studies), 54 samples of source rocks and bitumen were selected for GC analysis. Sixteen of these samples were analysed using the GC/MS technique after evaluation of the gas chromatograms (checking for 'biomarkers' in the C_{27} to C_{32} range) (Østfeldt, 1987b). About 115 samples from Peary Land had been analysed prior to the initiation of the Nordolie project (Rolle & Wrang, 1981). Many of these samples are, however, postmature and lean in organic material and provide only little information on the petroleum geology of the region.

Crushed samples (50–100 g except for the pure bitumens) were extracted with methylene chloride for 24 h in a Soxhlett extraction apparatus. The extract was filtered, the solvent removed by evaporation, and the amount of extract determined.

The extract (50–100 mg) was separated on an open silica column by stepwise elution with *n*-hexane, methylene chloride and methanol, yielding saturated, aromatic and polar (= hetero compounds) fractions, respectively. Asphaltenes are retained on the column. After evaporation of the solvent, the weight of the individual fractions was determined, and the relative distribution of saturate (sa), aromatic (ar) and polar (po) compounds was calculated.

GC analyses of the total saturate fraction were carried out on a Hewlett Packard 5840 gas chromatograph, fitted with a 25 m × 0.3 mm (int. diam.) fused silica capillary column coated with a cross-linked methyl silicone stationary phase. The column was operated from 80° to 300°C at 6°C/min. and the effluent detected with a FID detector. From the gas chromatogram, the pristane

to phytane (Pr/Ph) and pristane to C_{17} n -alkane (Pr/ nC_{17}) ratios were calculated (see fig. 11).

GC/MS analyses were carried out using a Finnigan 1020 GC/MS system connected with an on-column injector onto a 25 m \times 0.22 mm (int. diam.) fused silica capillary column coated with cross-linked methyl silicone phase. The column was operated from 70° to 310°C at 5°C/min. The column led directly into the ion source of the mass spectrometer operating at an ionizing voltage of 70 eV.

The mass spectrometer was operated in the Multiple Ion Detection (MID) mode, scanning 10 ions (m/e 82, 123, 177, 183, 191, 205, 217, 218, 231, and 259) every second. From the integrated ion-chromatograms of m/e 191 (triterpanes), 217 and 218 (steranes and diasteranes), several source, lithology and maturity-dependent parameters were calculated (see Østfeldt, 1987a,b for details). The other seven ion-chromatograms (not integrated) were used to support the tentative peak assignment.

Reflectance and fluorescence studies

Reflectance and fluorescence measurements of graptolites, kerogen and bitumen were carried out at the source rock laboratories of DGU and GGU. The graptolite-rich samples were collected for initiation of a research programme on graptolite reflectance (Stouge *et al.*, 1988) and about 45 samples were prepared shortly after the two field seasons. About 75 samples were selected for reflectance and fluorescence measurements of kerogen and bitumen after initial evaluation of screening data (Thomsen & Guvad, 1987). Only samples with relatively high TOC values or visible fluorescence in the palynological slides were considered further.

The investigations were carried out on whole rock samples, either crushed to a grain size of 2 to 3 mm or prepared as rock fragments orientated perpendicular to the lamination. The samples were embedded in a cold setting epoxy resin and then ground and polished using

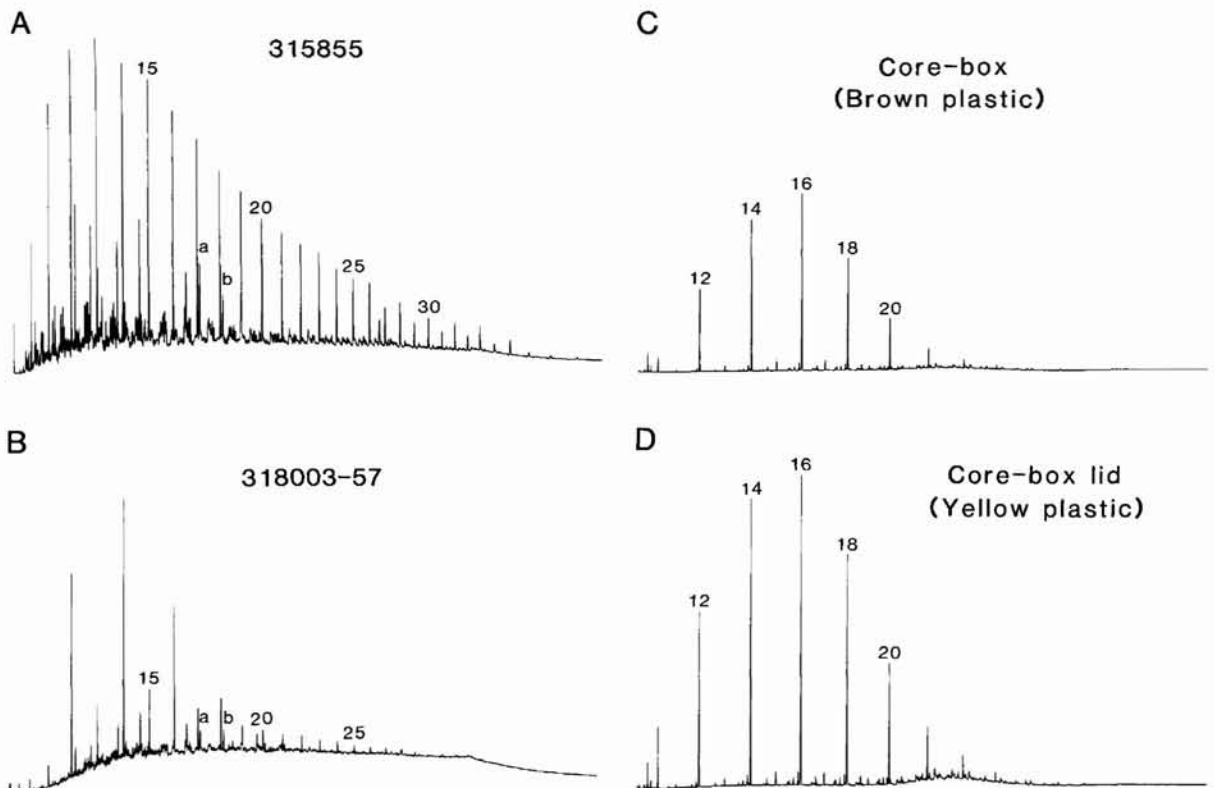


Fig. 11. Selected gas chromatograms of the saturated fraction. a: pristane, b: phytane, numbers are n -alkane carbon numbers. (A) Mature source rock. Sample 315855, Henson Gletscher Formation, Freuchen Land. (B) Polluted bitumen rock, note anomalously high values of C_{12} , C_{14} and C_{16} n -alkanes. Sample 318003-57, Henson Gletscher Formation, Freuchen Land. (C) Extract of core-box (brown plastic). Note the high contents of C_{12} , C_{14} , C_{16} , C_{18} and C_{20} n -alkanes. (D) Extract of core-box lid (yellow plastic). Note the composition similar to the core-box.

1/4 micron diamond powder for the final polish. Two microscope systems have been employed for measurement, in both cases using oil immersion: A reflected-light Zeiss photomicroscope equipped with an MP03 photometer and digital readout, using a 40× Epi-pol oil immersion objective, working with a plane glass reflector. Data are collected and processed by an MPS 3000 microcomputer. A Leitz MPV-SP reflected-light microscope equipped with a pol-opak illuminator, plane glass reflector and a 32× oil immersion objective. Data are collected and processed by an MPS 3000 microcomputer.

Illumination for both systems was through a green filter with peak transmission at 546 nm. The photomultipliers were calibrated against standards with reflectance values of 0.516 and 1.26% R_0 . Quantitative measurements of fluorescence were carried out using the Zeiss microscope system, fitted with a continuous filter monochromator and a 100W high-pressure mercury lamp connected to a stabilized power supply. For UV excitation the Zeiss F1 fluorescence reflected-light illuminator is used with a 25× neofluar objective.

Carbon isotope studies

Carbon isotope measurements of kerogen, bitumen and source rock extracts were carried out at the Stable Isotope Laboratory at the Geological Institute, University of Copenhagen. This method was brought into use in the later part of the project, mainly in order to correlate observed migrated bitumens with the two main source rock units but also to provide additional information on the depositional environment.

Measurements were carried out on both kerogen and extracts of the source rock ($n = 16$) and extracts of the migrated bitumen ($n = 25$). The total extract (see previous section on GC) was employed, typically from the same samples as used in the GC, GC/MS programme. Kerogen was separated from rock samples by decalcification with HCl followed by methylene chloride extraction to remove bitumen and dried at 90°C.

Combustion to CO_2 (of both carbon in kerogen and in extracts) was carried out in an oxygen-helium atmosphere at 900°C with copper oxide as catalyst. The evolved gas was purified over copper and silver at 600°C and transferred to a Finnigan MAT 250 triple collector mass spectrometer. The ratio between ^{13}C and ^{12}C is reported as per mille deviations from the PDB-standard using the δ -function (Epstein *et al.*, 1951). Reproducibility is better than 0.05‰ on the δ -scale.

X-ray diffraction of kerogen

X-ray diffraction of kerogen concentrates was carried out at the Laboratory of Clay Mineralogy, DGU (Koch, 1987). This method, rarely applied in source rock studies, was introduced in the final stage of the working programme with the special aim of characterizing and ranking thermally postmature samples.

Fifteen organic-rich samples were prepared, eight of these represent a profile through the Cambrian-Ordovician outer shelf and slope sequence with a known systematic increase in thermal maturity. The remaining Silurian samples have a known, but highly scattered, thermal maturity.

Depending on the content of organic carbon 10–25 g of powdered sample was treated with 15% HCl at a temperature of 80°C (see details in Koch, 1987). Following centrifugation, washing with distilled water and drying, the solid material was treated with concentrated HF at 80°C for two days and finally dried.

The kerogen concentrate was investigated by X-ray diffractometry using a Phillips 1050 vertical goniometer equipped with a graphite monochromator in the diffracted beam and using $CoK\alpha$ as radiation. In addition to the kerogen study a semiquantitative evaluation of the mineral content of the samples was made. The X-ray diffractogram in fig. 12 illustrates the procedure for determination of the two parameters d_{002} and WHH_{002} (position and width at half height of the 002 'graphite peak'). A number of other peaks show the presence of mineral phases (either relicts or minerals formed during preparation).

Fission track studies

Fission track studies of apatite and zircon from sandstones were carried out at the Geological Institute, University of Copenhagen. This method was included in the later part of the 'Nordolie' project in order to obtain chronological information on thermal episodes and history of uplift.

Fifteen samples representing two cross sections, one in the area where the Henson Gletscher Formation outcrops and one in the Silurian shales in Nyeboe Land, were prepared. Sample material of 0.5 to 1.5 kg was crushed and separated by magnetic and heavy liquid methods (see details in Hansen, 1988). The separates were mounted in epoxy or teflon, polished, and etched to reveal spontaneous tracks.

Age determinations were carried out using the external detector method (Gleadow & Lowering, 1978) with the Fish Canyon and Mt. Dromedary apatites and zircons as age standards following the zeta calibration

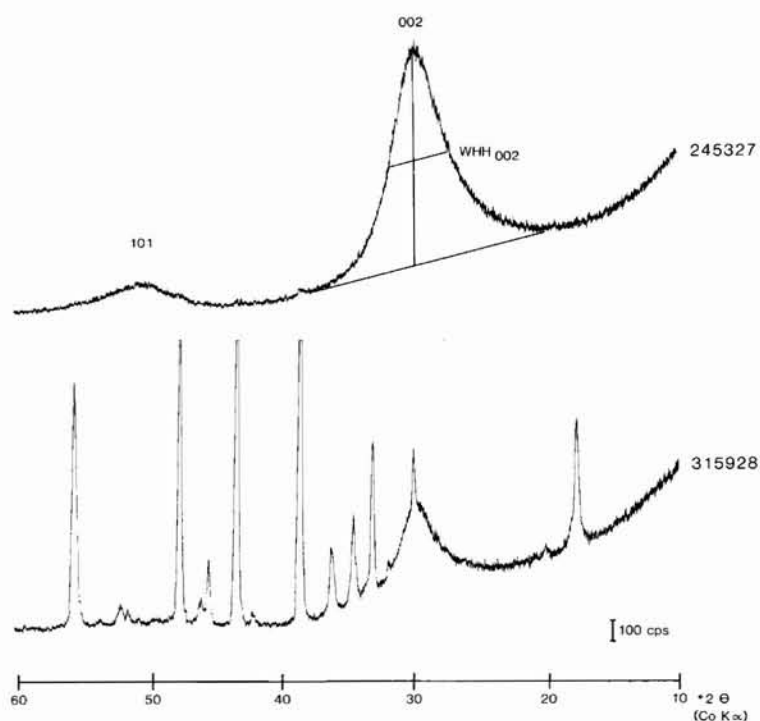


Fig. 12. X-ray diffractograms of samples 245327 and 315928 showing the procedure for determination of d_{002} and WHH_{002} . The sharp peaks indicate the presence of pyrite and fluorides formed during preparation.

procedure described by Hurford & Green (1983). NBS SRM 612 and Corning CN1 and CN2 glasses were employed for monitoring the neutron fluence. The polished and etched apatites and zircons were irradiated together with high quality low-uranium mica detectors at the J1 facility of the HERALD reactor in Aldermaston, U.K., or at the Risø National Laboratory, Denmark.

All measurements for calculating the fission track length distribution in apatite were carried out on horizontal, confined tracks as suggested by Gleadow *et al.* (1986).

Fluid inclusion studies

Studies of fluid inclusions and stable isotopes of diagenetic calcite and dolomite associated with bitumen were carried out at the Geological Institute, Copenhagen University. These methods were employed in the later part of the project in order to obtain information on the temperature and chemistry of pore waters at the time of diagenetic growth and migration of hydrocarbons (Jensenius, 1987).

Only samples from four localities, all with macroscopically identified bitumen, were investigated.

Fluid inclusion microthermometry was performed with the aid of a Chaimeca heating and freezing stage (Poty *et al.*, 1976) calibrated from 96°C to 400°C. The

stable isotopes were analysed by the following procedure at the Stable Isotope Laboratory, Geological Institute, University of Copenhagen. Approximately 30 mg of carbonate were crushed to a grain size between 64 and 150 μm , treated with sodium hypochlorite to remove migrated hydrocarbons and kerogen, followed by reaction in vacuum with concentrated H_3PO_4 . The evolved CO_2 was cleaned over dry ice isopropanol mixture and subsequently admitted to the Finnigan Mat-250 isotope mass spectrometer.

Porosity/permeability

Measurements of the petrophysical parameters (porosity, permeability and grain density) were carried out commercially by the Core Analysis Laboratory of DGU. Approximately 60 orientated plugs (25 mm \times 25 mm), mostly from Cambrian sandstones, were analysed. Following drying, the specific permeability is measured by flowing nitrogen through the plug. The porosity and grain density is determined applying a double chambered helium porosimeter and a picometer with calibrated mercury pump (Springer, 1987).

Thin sections of all analysed samples from the plugs or nearby slabs have been stained for carbonate identification by Dickson's method (Allman & Lawrence, 1972) and studied in detail.

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Plate 1. Kerogen

- A. Sample with a relatively low (0.21% TOC) kerogen content and a dominance of finely disseminated amorphous kerogen in a silica gel, Lafayette Bugt Formation, Nyeboe Land, GGU 316490-1, unsieved organic material.
- B. As A., Lafayette Bugt Formation, Nyeboe Land, GGU 316490-2, sieved organic material (on 10 μm nylon mesh).
- C. Sample with a relatively moderate (1.15% TOC) kerogen content and small to moderate amounts of large amorphous kerogen particles, Thors Fjord Member, Nares Land, GGU 318007-18-1 unsieved organic material.

- D. As C., Thors Fjord Member, Nares Land, GGU 318007-18-2, sieved organic material (on 10 μm nylon mesh).
- E. Sample with a relatively large (5.09% TOC) kerogen content and a dominance of large amorphous kerogen particles, Thors Fjord Member, Nares Land, GGU 318007-32-1, unsieved organic material.
- F. As E., Thors Fjord Member, Nares Land, GGU 318007-32-2, sieved organic material (on 10 μm nylon mesh).

Scale bar: 20 μm .

Plate 2. Cambrian palynomorphs

- A. Acritarch-like folded alga. Middle Cambrian Sydpasset Formation, Freuchen Land, MGUH 19334 from GGU 315873-2; 139.5-13.9.
- B. Acritarch-like folded alga. Middle Cambrian Sydpasset Formation, Freuchen Land, MGUH 19335 from GGU 315873-2; 150.6-7.0.
- C. Two acritarch-like folded algae. Middle Cambrian Sydpasset Formation, Freuchen Land, MGUH 19336 (large light body), MGUH 19337 (dark small body), both from GGU 315873-2; 122.0-15.3.
- D. Acritarch-like folded alga. Middle Cambrian, Ekspedition Bræ Formation, Freuchen Land, MGUH 19338 from GGU 324217-2; 139.2-14.9.
- E. Acritarch-like folded alga. Middle Cambrian, Ekspedition Bræ Formation, Freuchen Land, MGUH 19339 from GGU 324300-2; 131.2-13.4.

- F. Acritarch-like folded alga. Middle Cambrian, Ekspedition Bræ Formation, Freuchen Land, MGUH 19340 from GGU 324217-2; 141.9-11.4.
- G. Lump of algal or spore-like elements. Middle Cambrian, Ekspedition Bræ Formation, Freuchen Land, MGUH 19341 from GGU 324300-2; 127.8-14.4.
- H. Diad-like lump of algal or spore-like elements. Middle Cambrian, Ekspedition Bræ Formation, Freuchen Land, MGUH 19342 from GGU 314300-2; 138.5-17.8.
- I. Lump of alga or spore-like elements. Middle Cambrian, Ekspedition Bræ Formation, Freuchen Land, MGUH 19343 from GGU 324300-2; 157.7-14.7.

Scale bar: 20 μm .

Plate 3. Ordovician palynomorphs

- A. Acritarch. Upper Ordovician Troedsson Cliff Member, Washington Land, MGUH 19344 from GGU 316968-2; 145.3-17.8.
- B. Acritarch. Upper Ordovician - Lower Silurian Aleqatsiaq Fjord Formation, Washington Land, MGUH 19345 from GGU 316085-4; 124.1-21.3.
- C. Graptolite fragment, Upper Ordovician Troedsson Cliff Member, Washington Land, MGUH 19346 from GGU 316968-2; 128.1-2.9.
- D. Scolecodont, Upper Ordovician, Troedsson Cliff Member, Washington Land, MGUH 19347 from GGU 316968-2; 148.8-15.1.
- E. Alga. Upper Ordovician - Lower Silurian Aleqatsiaq Formation, Nyeboe Land, MGUH 19348 from GGU 316103-2; 135.0-15.0.
- F. Filamentous alga. Upper Ordovician - Lower Silurian Aleqatsiaq Formation, Washington Land, MGUH 19349 from GGU 316058-2; 135.1-4.2.

- G.-L. Spores with trilete rays. Upper Ordovician, Troedsson Cliff Member, Washington Land (Nøhr-Hansen & Koppellus, 1988).

G.-I. *Besselia nunaatica*, MGUH 17539 from GGU 316968-2; 125.5-8.3.

G. Distal view illustrating the minute ornamentation.

H. Equatorial view.

I. Internal proximal view.

J. *Besselia nunaatica*, two connected spores, internal proximal view, MGUH 17541 from GGU 316968-2; 155.1-11.9.

K.-L. *Besselia nunaatica*, MGUH 17542 from GGU 316968-2; 123.8-15.9.

K. Distal view illustrating the ornamentation.

L. Internal proximal view.

Scale bar: 20 μm .

Plate 4. Silurian palynomorphs

- A. Chitinozoan, *Angochitina* cf. *A. elongata*. Upper Silurian Wulff Land Formation, Wulff Land, MGUH 19350 from GGU 315950-3; 136.9-17.2.
- B. Chitinozoans, *Linochitina erratica*. Upper Silurian Wulff Land Formation, Wulff Land, MGUH 19351 from GGU 315950-2; 154.3-9.6.
- C. *Retiolites*, graptolite fragment. Upper Silurian, Wulff Land Formation, Wulff Land, MGUH 19352 from GGU 315950-3; 155.1-11.1.
- D. Graptolite fragment, Upper Silurian, Wulff Land Formation, Wulff Land, MGUH 19353 from GGU 315950-2; 127.3-5.7.
- E.-H. Trilete spore-like bodies, figs E and F with a degraded bitumen-like appearance.
- E. Lower Silurian Lafayette Bugt Formation, Washington Land, MGUH 19354 from GGU 211760-2; 143.3-17.2.
- F. Upper Silurian Wulff Land Formation, Wulff Land, MGUH 19355 from GGU 315950-3; 15950-3; 155.5-8.2.

G. Upper Silurian Nyeboe Land Formation, Nyeboe Land, MGUH 19356 from GGU 319234-2; 119.3-11.0.

H. Upper Silurian Nyeboe Land Formation, Wulff Land, MGUH 19357 from GGU 319210-3; 130.6-21.4.

I. Spherical folded algae, acritarchs? Lower Silurian Lafayette Bugt Formation, Hall Land, MGUH 19358 from GGU 324157-2; 144.2-8.5.

J. Tubular structure. Upper Silurian Nyeboe Land Formation, Wulff Land, MGUH 19359 from GGU 319210-3; 146.6-16.5.

K. Tubular structure. Upper Silurian Nyeboe Land Formation, Nyeboe Land, MGUH 19360 from GGU 319234-2; 138.8-8.0.

L. Rounded drop-shaped palynomorphs. Lower Silurian Lafayette Bugt Formation, Washington Land, MGUH 19361 from GGU 316061-2; 137.1-14.8.

Scale bar: 20 μm .

Plate 1. Kerogen

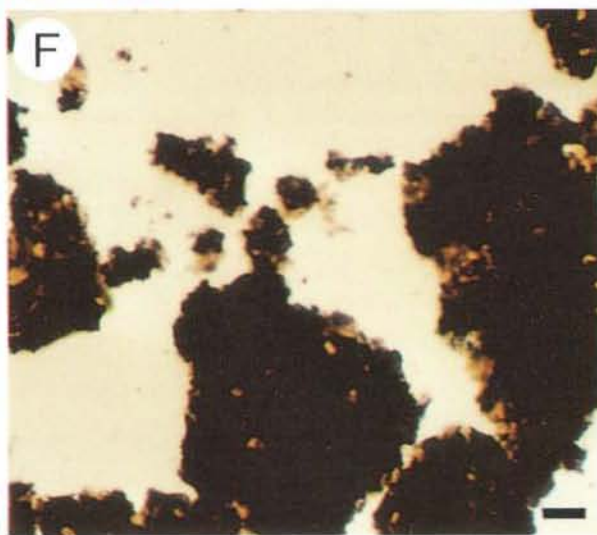
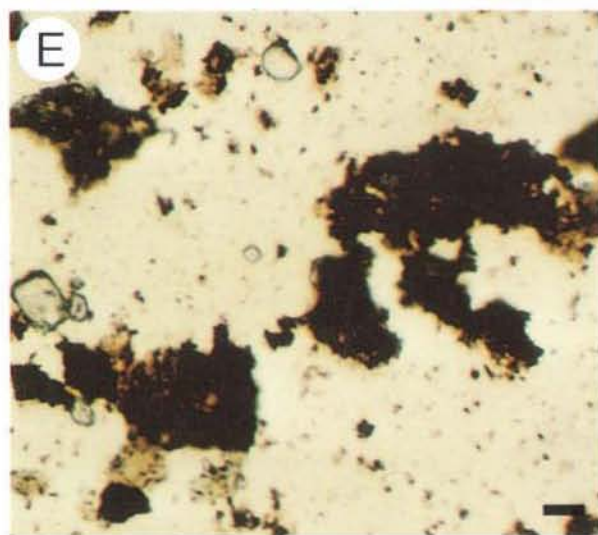
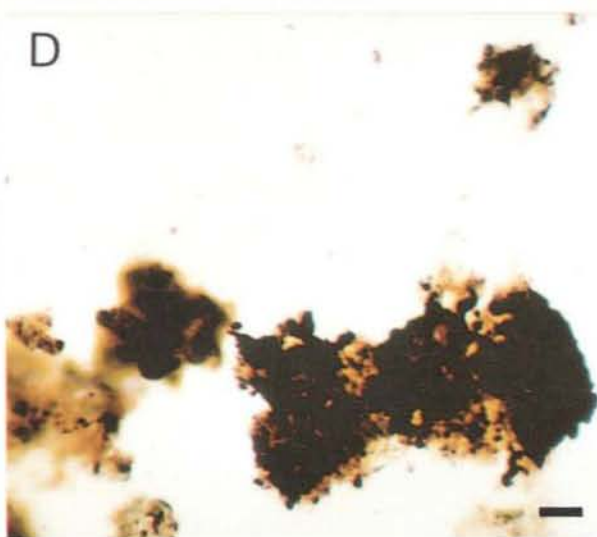
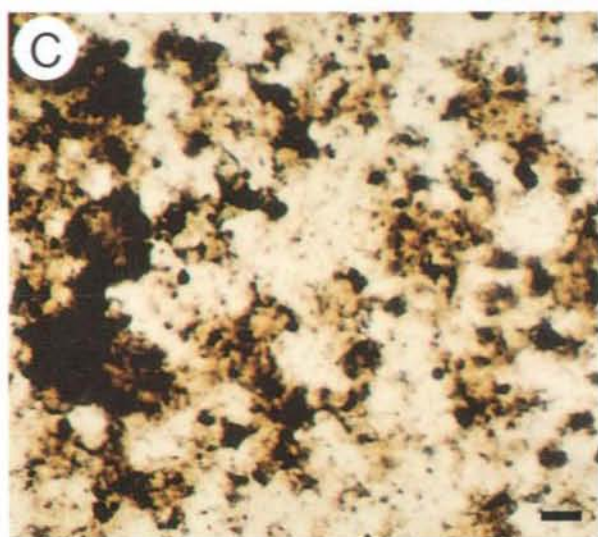
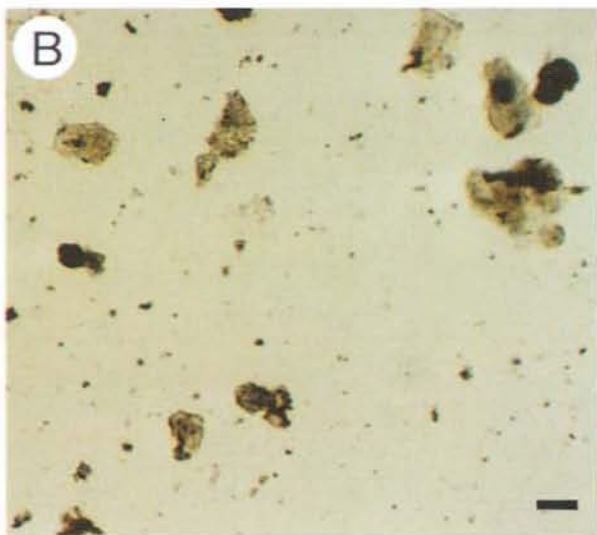
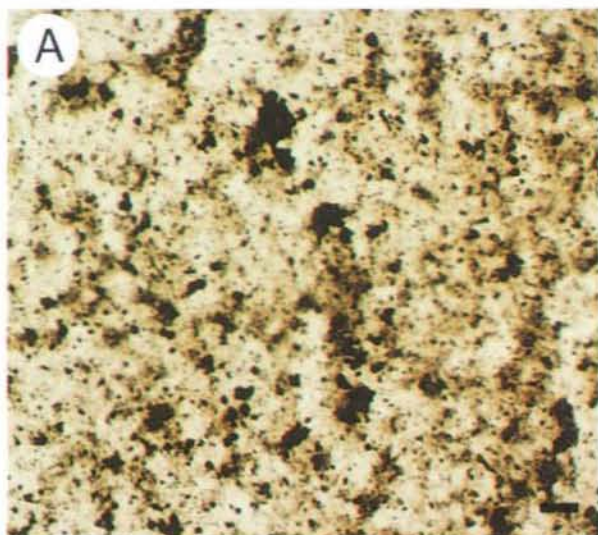


Plate 2. Cambrian palynomorphs



Plate 3. Ordovician palynomorphs

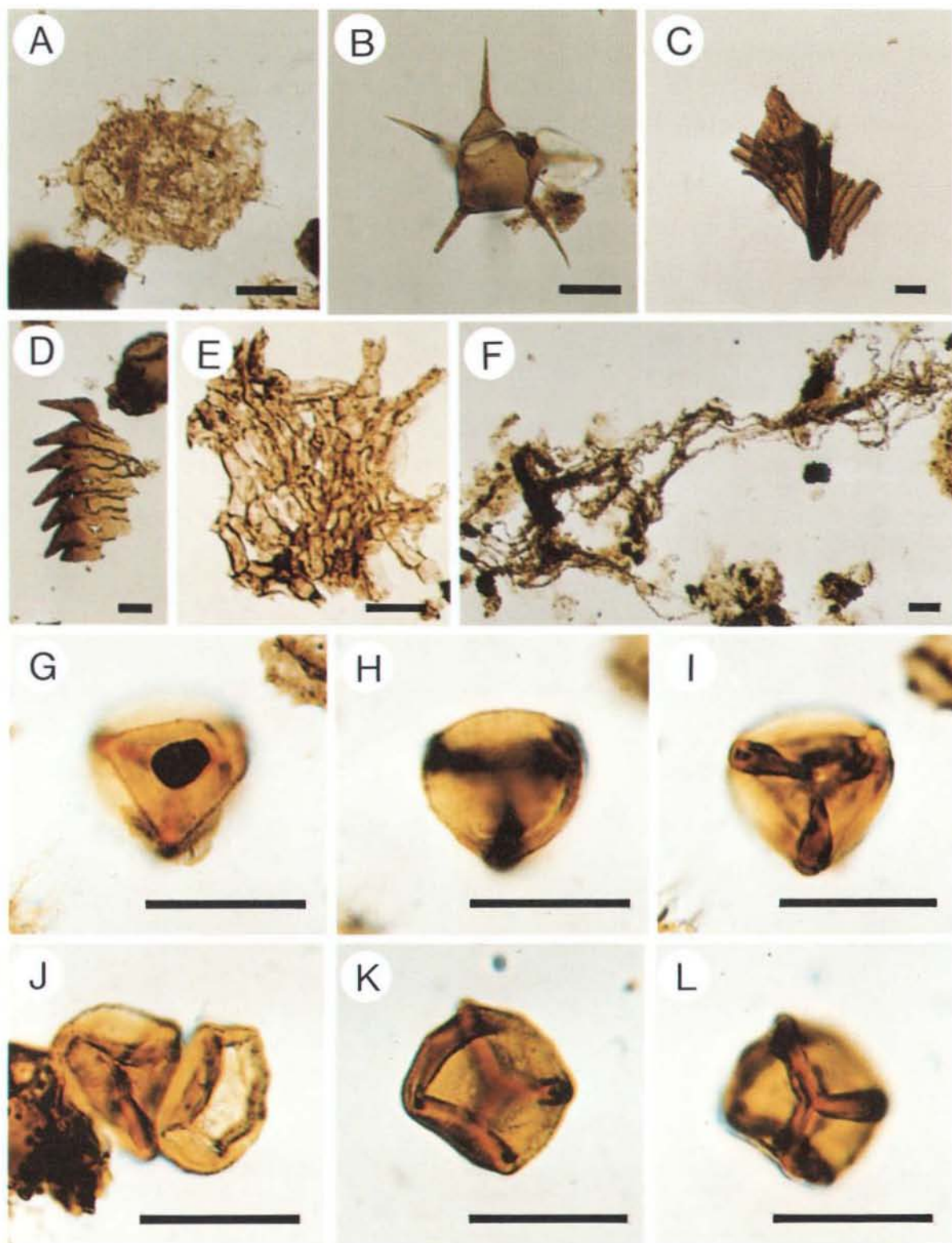


Plate 4. Silurian palynomorphs

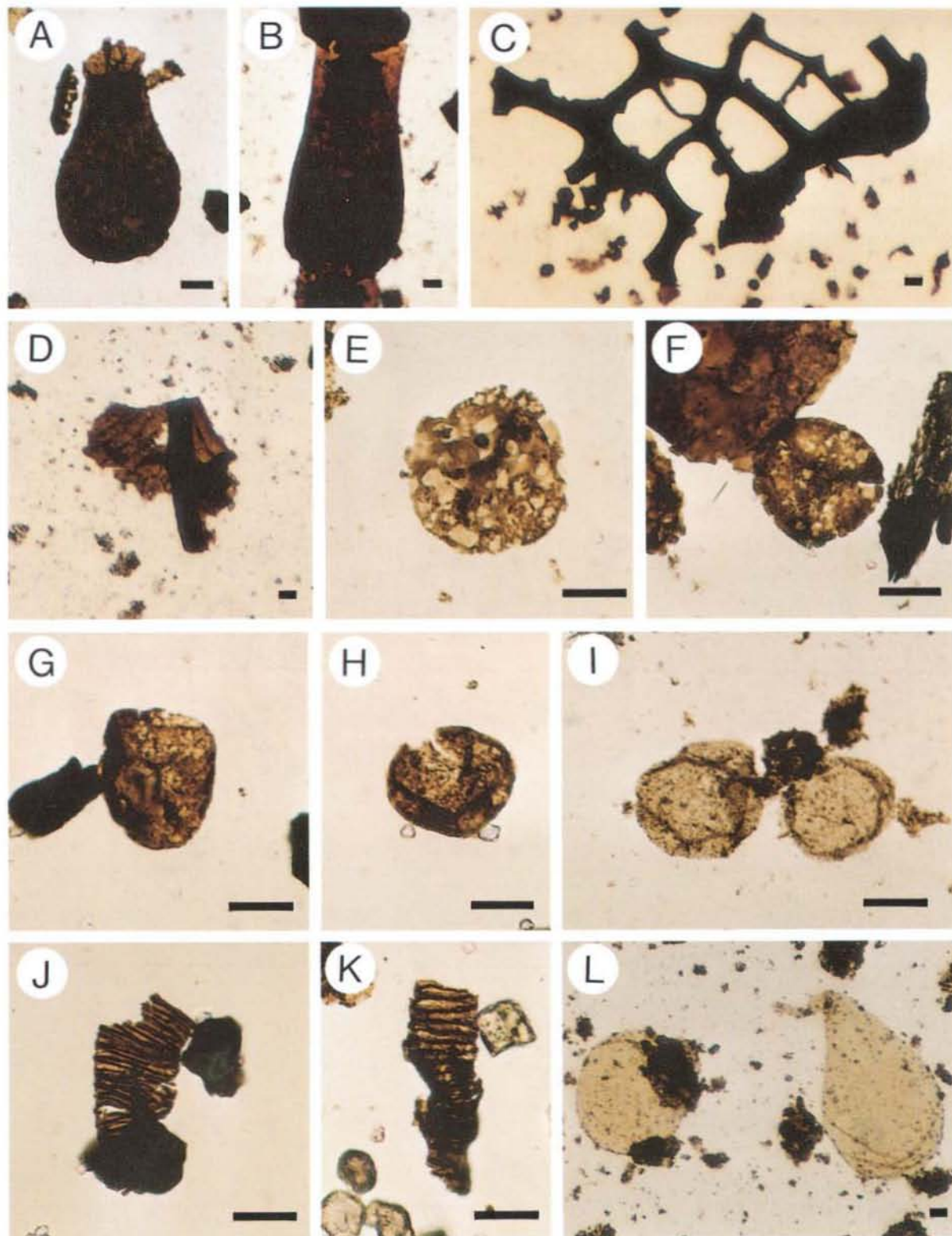


Plate 5. Progressive coloration of amorphous kerogen with increasing thermal alteration

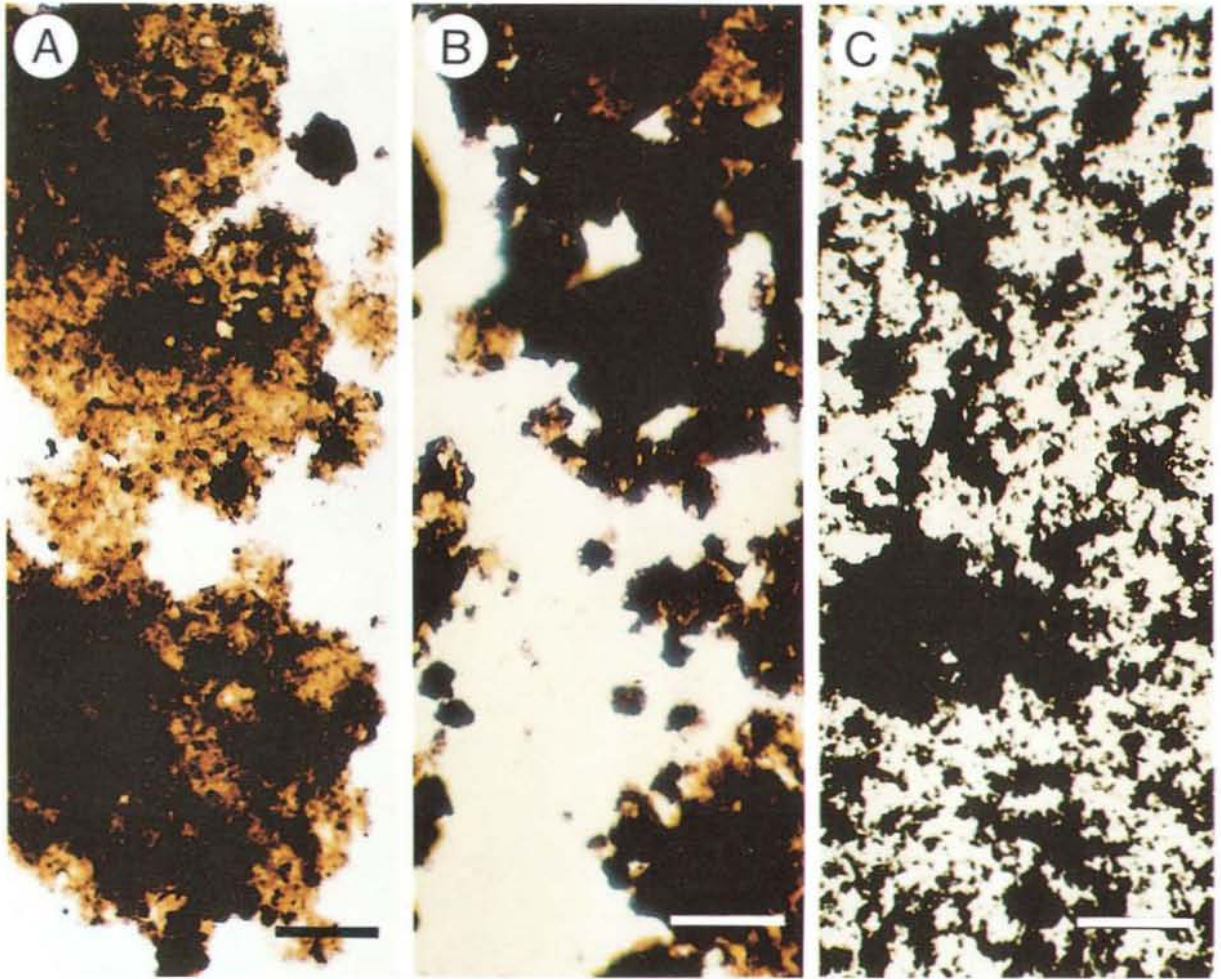


Plate 5. Progressive coloration of amorphous kerogen with increasing thermal alteration

Scale bar: 50 μm

A. TAI: (2)-2⁺, GGU 211759-2. B. TAI: 2⁺-(3-), GGU 324405-2. C. TAI: 4⁺, GGU 316475-1.

Plate 6. Change in structure of amorphous kerogen with increasing thermal alteration as observed in the scanning electron microscope

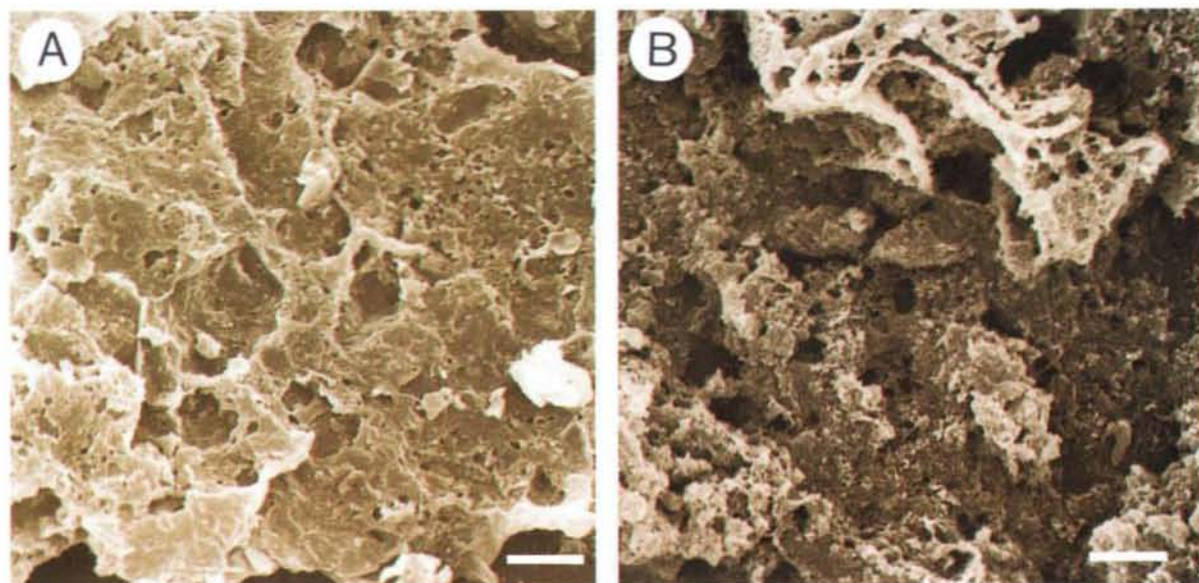


Plate 6. Change in structure of amorphous kerogen with increasing thermal alteration as observed in the scanning electron microscope

A. TAI: 2⁺-(3), T_{max} : 446, GGU 324405-2, scale bar: 10 μ m.

B. TAI: 4⁺, T_{max} : n.d., GGU 316475-2, scale bar: 10 μ m.

Plate 7. Field appearance of bitumen

A. Seeping asphalt from southern Wulff Land (equivalent to GGU 324200).

B. Hard solid bitumen in dolomite vug in the Sydpasset Formation (equivalent to GGU 324287-324299, core GGU 318003).

Plate 8. Macroscopic bitumen in slabs

A. Asphalt from seep in dolomite breccia. GGU 324200A.
B. Like A. Stained. D0, D1, D2, C1, C2 correspond to generations of dolomite and calcite.

C. Asphalt from seep in dolomite breccia. Stained. Same generations of carbonates as A and B. GGU 324200E.

D. Hard solid bitumen in vugs and veins in dolomite grainstone. DO: dolomite grainstone, D1: saddle dolomite. Core GGU 318003.

Plate 9. Bitumen in thin section

A. Bitumen in coral. GGU 324130B, Lafayette Bugt Formation, Nyeboe Land. Plane light, stained, C: calcite, Fe-C: Fe-rich calcite, B: bitumen, scale bar: 1 mm.
B. Bitumen in coral. GGU 316067, Lafayette Bugt Formation, Washington Land. Crossed nicols + gypsum plate. Q: quartz, C: calcite, B: bitumen, scale bar: 1 mm.
C. Bitumen-filled fracture in calcarenite. GGU 318013-09, Lafayette Bugt Formation, Nyeboe Land. Plane light, scale bar: 2 mm.

D. Two-phased bitumen (black and yellow) (B1, B2) in saddle dolomite veins (D1) cross-cutting dolomite grainstone (D0). See close-up (arrow) in Plate 11. GGU 318003-53, Henson Gletscher Formation, Freuchen Land. Plane light, scale bar: 2 mm.

E.-F. Saddle dolomite vein (D1) in dolomite grainstone (D0). Bitumen occurs as impregnation in D0 (B1), as residual matter in the contact between D0 and D1 (B2, R_o : 0.92%) and in the centre of the vein (B3, R_o : 1.21%). GGU 318003-21, Sydpasset Formation, Freuchen Land. Plane light, scale bars: 1 mm.

Plate 10. Bitumen in palynologically prepared samples observed in microscope or in SEM

A. Bitumen with flaky appearance (note crystal impressions). GGU 315172-1, Ryder Gletscher Group Fm 6, Wulff Land. Scale bar: 25 μ m.
B. Globular bitumen. GGU 315865-2, Aftenstjernesø Formation, Nares Land, scale bar: 50 μ m.
C. Globular bitumen which has been extruded during and after sample preparation by the xylene-containing mounting medium. GGU 315199, Ryder Gletscher Group Fm 6, Warming Land. Scale bar: 12.5 μ m.

D. Bitumen with flaky appearance (note crystal impressions). GGU 315172-1, Ryder Gletscher Group Fm 6, Wulff Land. Scale bar: 10 μ m.
E. Bitumen mirroring imprints of crystals from coral space. GGU 316067-2, Lafayette Bugt Formation, Washington Land. Scale bar: 10 μ m.
F. Thread-like bitumen. GGU 324453, Buen Formation, Wulff Land. Scale bar: 25 μ m.

Plate 7. Field appearance of bitumen



Plate 8. Macroscopic bitumen in slabs

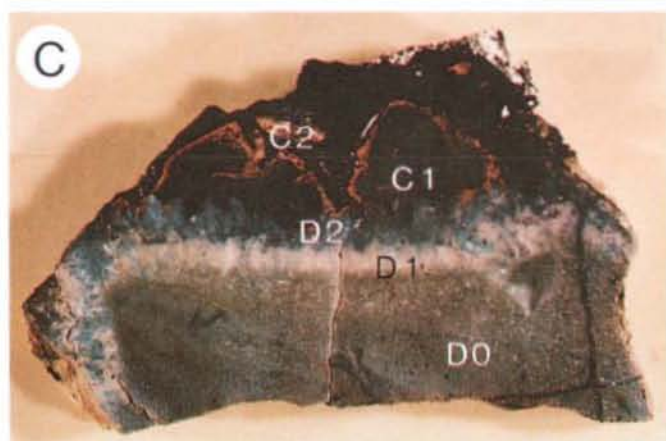
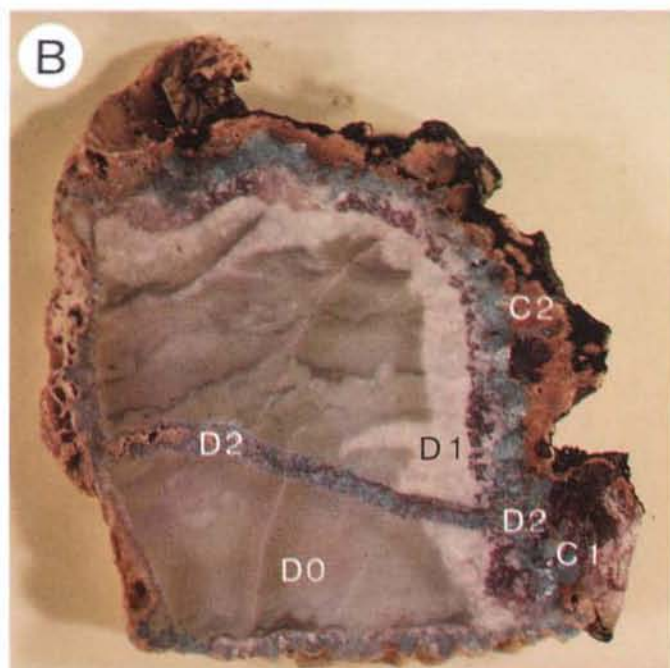
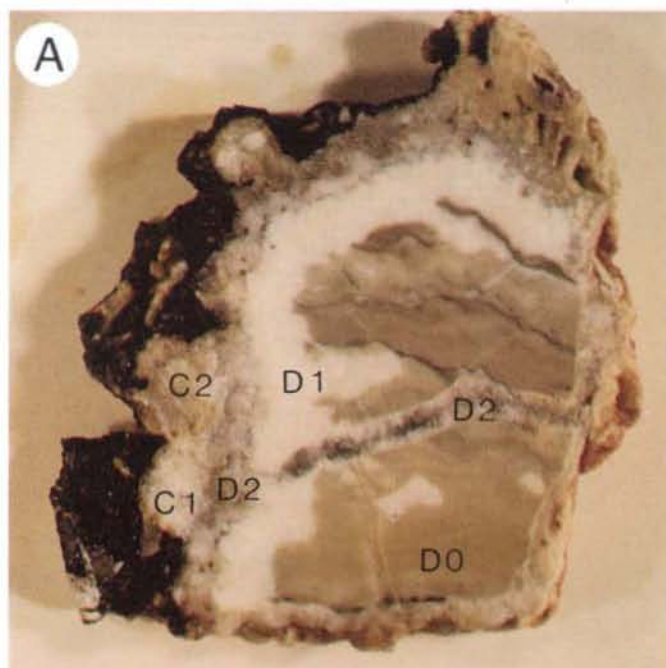


Plate 9. Bitumen in thin section

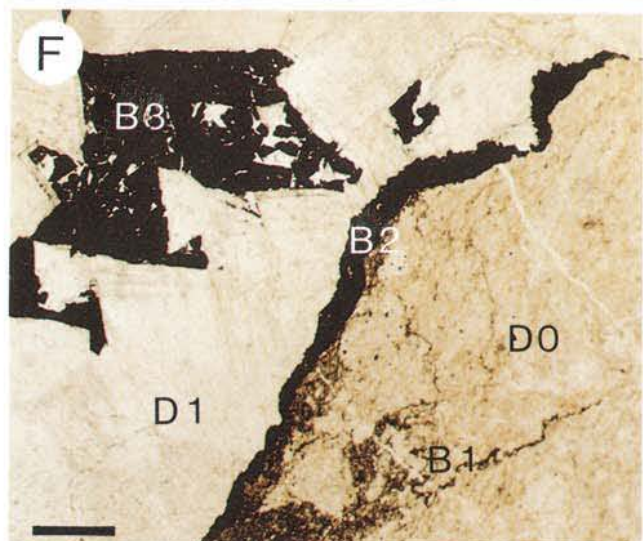
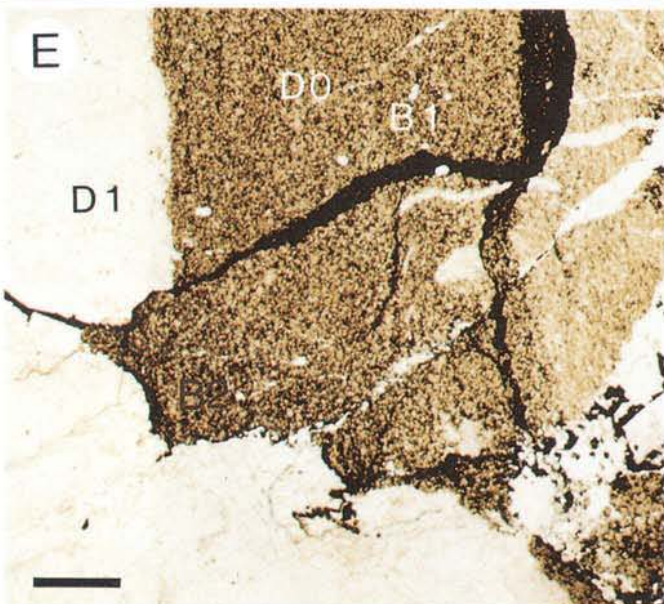
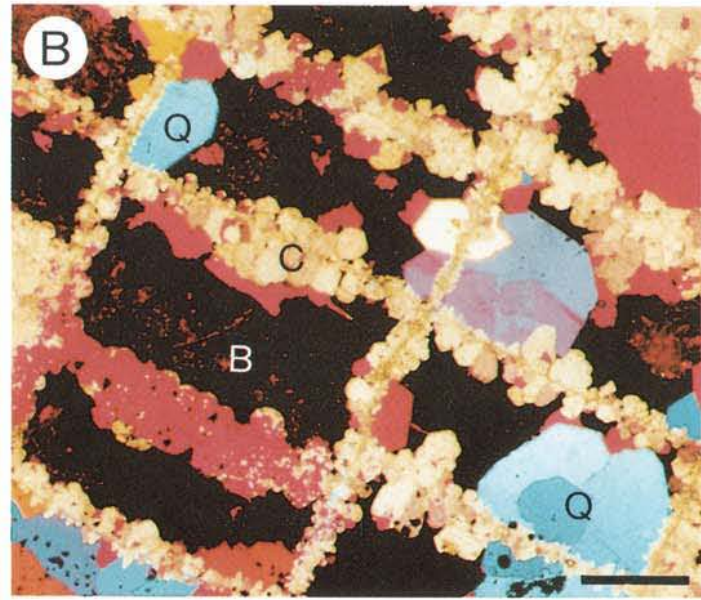
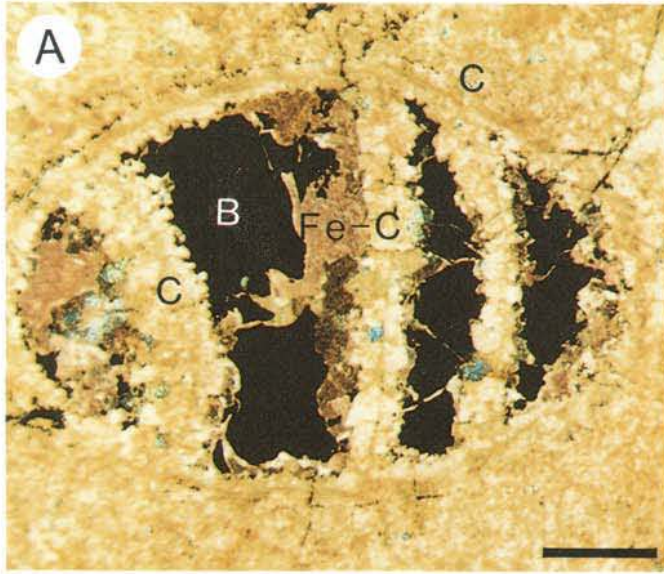


Plate 10. Bitumen in palynologically prepared samples observed in microscope or in SEM

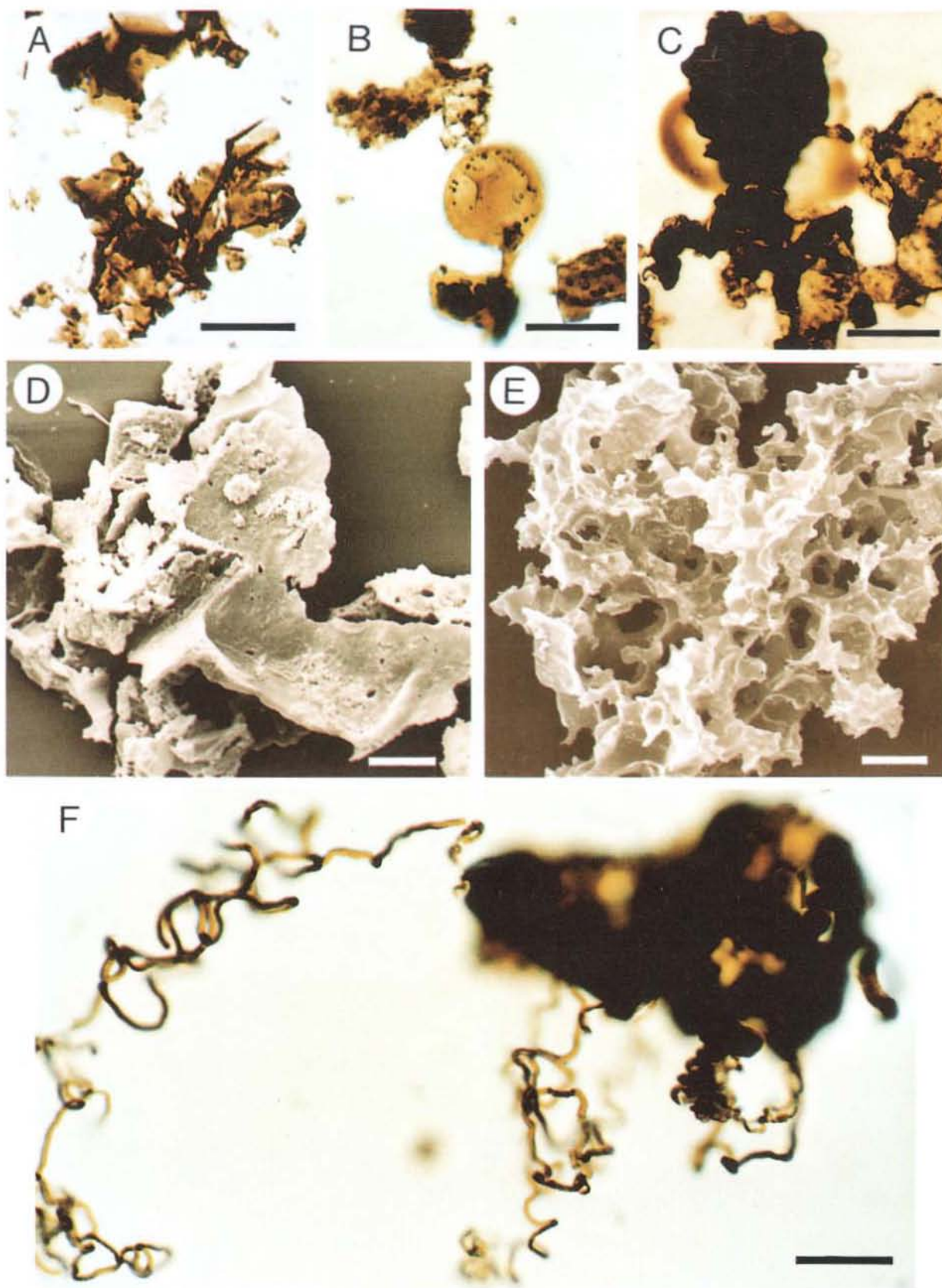


Plate 11. Bitumen in polished section

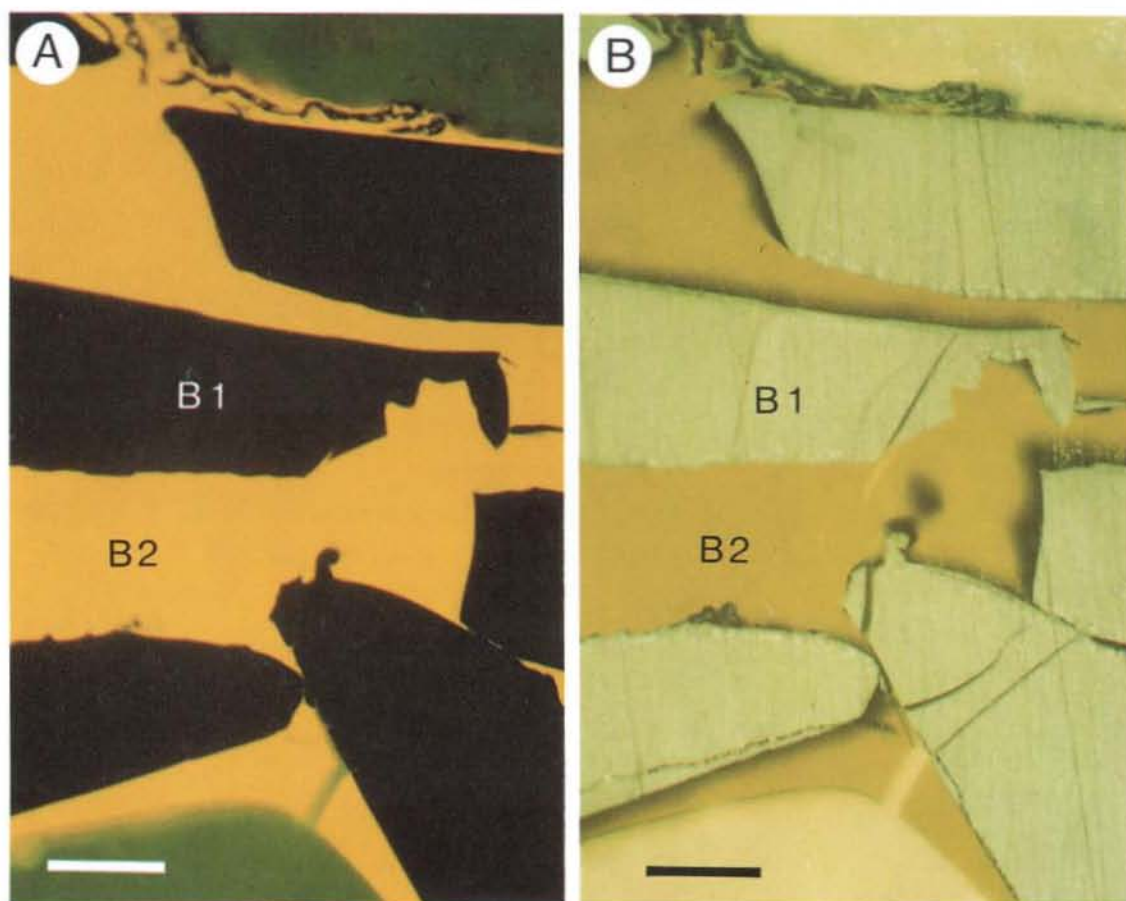


Plate 11. Bitumen in polished section

GGU 318003-53, Henson Gletscher Formation, Freuchen Land.

Scale bar: 50 μm

A. Fluorescent light photograph of two-phase bitumen.

B. Normal reflected light photograph of same field. The yellow-fluorescent low-reflecting bitumen (B2) has a R_o of 0.08% and the dark non-fluorescent high-reflecting bitumen (B1) a R_o of 1.17%.

Cover picture

Cambrian and Ordovician strata at Blue Cliffs, Wulff Land, North Greenland. Photo: J. Lautrup, GGU.