

SOME SUGGESTIONS ON POLLEN AND SPORE DIAGNOSES

G. ERDTMAN

Palynological Laboratory, Stockholm-Solna, Sweden

ABSTRACT—The author discusses the problem of discrepancies in terminology and offers suggestions concerning terminological unification in the case of pollen and spores.

INTRODUCTION

MANY new and important botanical and geological facts have accumulated as a consequence of the increasing activity in fossil pollen and spore research. It is, however, unfortunate that the bright outlook of this branch of micro-palaeontology is somewhat marred by the fact that descriptions of the pollen or spore "species" are sometimes rather vague or even imperfect to such an extent that the value of the



"species" can rightly be contested. There are probably several reasons for this. A micro-palaeontologist without propaedeutical training in the morphology of recent pollen grains and spores will thus hardly realise the great factual possibilities of submitting detailed pollen and spore diagnoses based on symmetry, apertures, shape, sporoderm stratification, etc. Contributory reasons are probably to be found in still prevailing terminological discrepancies, as well as in the fact that, owing to technical difficulties or lack of knowledge, it has not yet been possible to establish a sufficiently detailed and consistent terminology. Thus, pending some penetrating morphological analyses, the term "os", i.e., "the inner part of a composite aperture" should be regarded as a preliminary, collective term for a probably more or less heterogeneous group of morphological features [cf. e. g. the ora in the "colporate" grains in *Mendoncia aspera* (Erdtman 1952, fig. 6D, p. 31), *Geissoloma marginatum* (l. c., fig. 107, p. 183), *Myoporum laetum* (l. c., fig. 161,

p. 277), *Penaea mucronata* (l. c., fig. 184A, p. 317), *Rhoiptelea chiliantha* (l. c. Fig. 222, p. 380), *Erythrochiton brasiliensis* (l. c., fig. 226A, p. 389), *Berrya cordifolia* (l. c. fig. 248C, p. 435), and *Leea aequata* (l. c., fig. 257B, p. 451)].

In order to overcome at least some of the terminological discrepancies a few suggestions towards a terminological unification have recently been made (cf. Erdtman and Vishnu-Mittre 1957). In the present note this revised terminology is tentatively applied in a number of elementary pollen diagnoses accompanied by palynograms ($\times 1000$; from Erdtman 1952), picturing, in a more or less broad outline, the essential pollen-morphological characters.












A diagnosis should be sufficiently detailed to make it possible to draw a palynogram on the basis of the data given. Likewise, a palynogram does not fulfil its purpose if a satisfactory pollen or spore diagnosis cannot be based on it. Explicit and consistent diagnoses, preferably accompanied by palynograms, photomicrographs, electron micrographs, etc., will no doubt contribute towards a safer and better approach to the many interesting problems in fossil pollen and spore research.

DIAGNOSES

(for explanation of terms, see fig. 1 and the papers quoted in "References")

Peumus boldus (fig. 2; Monimiaceae; Chile, Werdermann 311): grains inaperturate, spherical, spinulose (diameter 33μ , spinules included).

Exine (spinules included) about 2μ thick, crassisexinous. Spinules about $1-1.5 \mu$, isometric, equidistant (about 2.5μ apart). Endosexine baculate (10).

Faegri and Iversen 1950 Iversen and Troels-Smith 1950	ERDTMAN 1952	POSITION OF APERTURES					Main subdivisions
		POLAR (proximal)	POLAR (distal)	ZONAL	GLOBAL	UNKNOWN	
	monolete						monolete
	trilete						trilete
	hilate						cataporate
colpate p.p.	sulcate						anacolpate
porate p.p.	ulcerate						anaporate
colpate p.p. stephanocolpate	colpate						zonicolpate
porate p.p. stephanoporate	porate						zoniporate
pericolpate	rugate						pancolpate
periporate	forate						panporate
colpate							colpate
porate							porate

Text-fig. 1—Aperture terminology. (From Erdtman and Vishnu-Mittre 1957.) Cf. also Addendum p. 52.

Hedycarya arborea (fig. 3; Monimiaceae; New Zealand): pollen grains united in \pm tetrahedral tetrads (diameter of tetrads 39μ), inaperturate or, more probably, anatenuate.

Exine about 1μ thick. Distal face with a diffuse lo-pattern (except at the supposed tenuitas and near the borderline between the individual grains).

Capsicodendron pimenteria (fig. 4; Canelaceae; Brazil, Hoehne 27074): grains anacolpate, biconvex, $27 (17+10) \times 34 \times 30 \mu$. Colpus about $22 \times 6 \mu$.

Exine about 2μ thick at equator. Sexine as thick as nexine or slightly thinner, smooth, not distinctly stratified.

N.B. The size figures comprise the length of the polar axis (27μ) and the length of the two equatorial diameters, the colpiferous (34μ) and the non-colpiferous (30μ). The figures in parenthesis ($17+10$) after the polar axis figures indicate that about 17μ of the polar axis falls within the colpiferous convexity and about 10μ within the non-colpiferous (cf. also Erdtman 1946).

Lemna gibba (fig. 5; Lemnaceae): grains anaportate, spinulose (diameter 22μ , spinules not included). Pore circular (diameter about 2μ).

Exine about 1.5μ thick (spinules not included). Spinules about 1.5μ long, \pm equidistantly spaced (about 4μ apart).

Oxalis acetosella (fig. 6; Oxalidaceae; Sweden, Erdtman s.n.): zonocolpate (3), subprolate ($53 \times 44 \mu$). Apocolpium diameter about 14μ . Colpi with pointed ends, about 1.5μ broad at equator, tenuimarginate.

Exine about 2.25μ thick. Sexine thicker than nexine, finely reticulate. Ectos-exinous part of muri thinner than endosexinous.

Myriophyllum spicatum (fig. 7; Haloragaceae; Sweden): grains zonocolpate (3-5), suboblate ($24 \times 28 \mu$), aspidote (diameter of aspides about 5.5μ). Colpi, when three, meridionally arranged; when four or five, with their axes biconvergent; short (according to Wodehouse 2.85 by 11.4μ).

Exine about 1μ thick. Sexine thinner than nexine. Colpi margins incrassate (due

to thickening of ectonexine; maximum thickness of colpi margins about 2.5μ).

N.B. By Faegri and Iversen (1950) the genus *Myriophyllum* is, (i.a.) mentioned in p. 146 under the class stephanoporate. According to their definitions and the illustrations in p. 131 (l.c.) the pollen grains in *M. spicatum* should be classified as 3-colpate (when provided with three meridional apertures) or as pericolpate (when provided with non-meridional apertures). According to Erdtman 1952, the grains are either 3-colpate or 4-5-rupate.

Cordia alliodora (fig. 8; Boraginaceae; Mexico, Gentle 1561): grains zonocolpate (3), suboblate (about $26 \times 30 \mu$). Apocolpium diameter about 21μ . Colpi about 2μ wide at equator, tenuimarginate. Ora lalongate (about $3 \times 11 \mu$).

Exine about 2μ thick (spinules not included). Sexine tectate (consisting of suprategillar spinules about 0.5μ long and 1.5μ apart, tegillum, and infrategillar bacula), approximately as thick (spinules not included) as nexine.

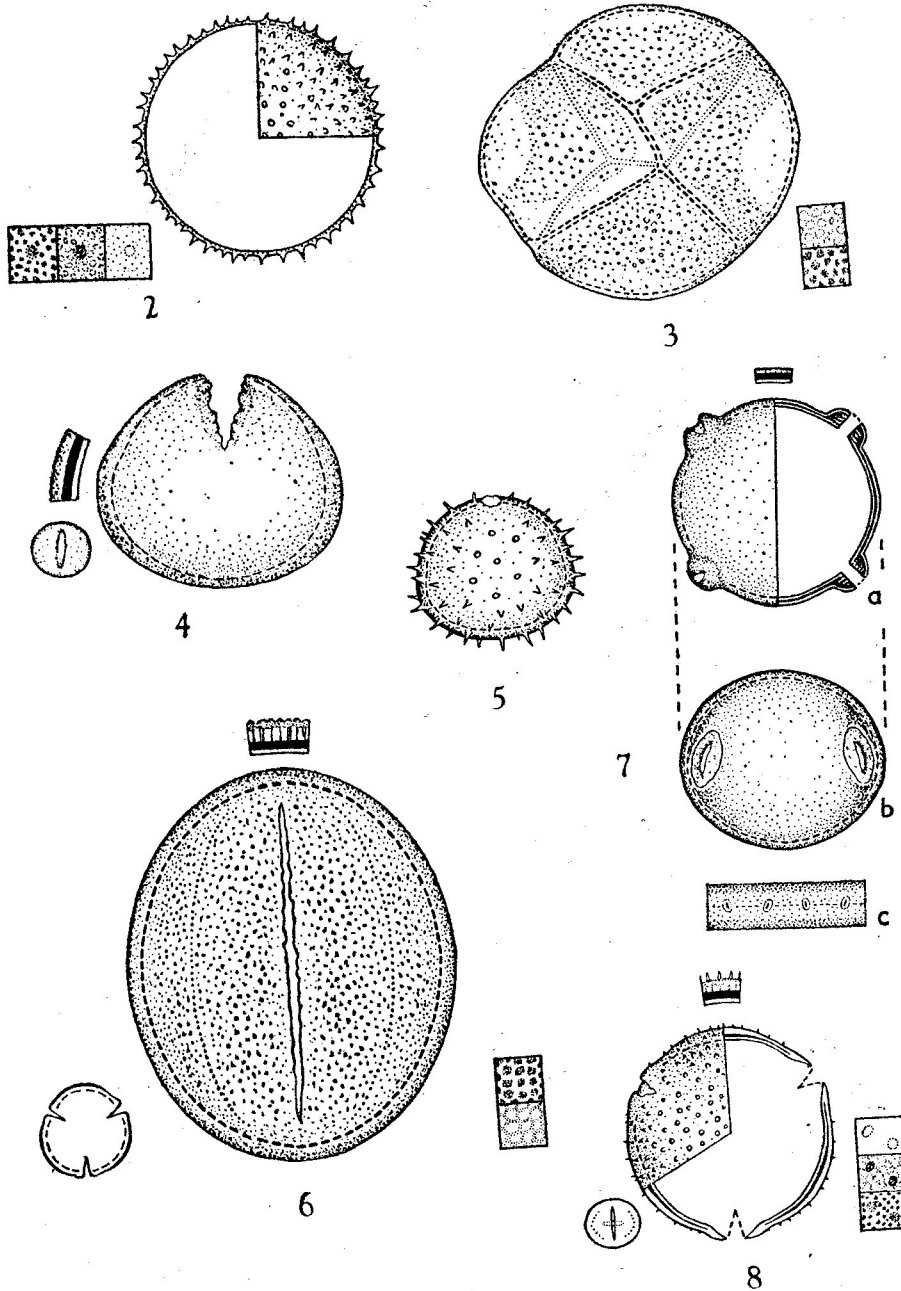
Franklandia fucifolia (fig. 9; W. Australia; Harvey s.n.): grains zonoporate (3), oblate spheroidal ($82 \times 92 \mu$). Pores circular (diameter about 10μ).

Exine about 14μ thick. Sexine consisting of densely spaced blunt rods (about 7μ long by 2μ broad). Nexine about 7μ thick.

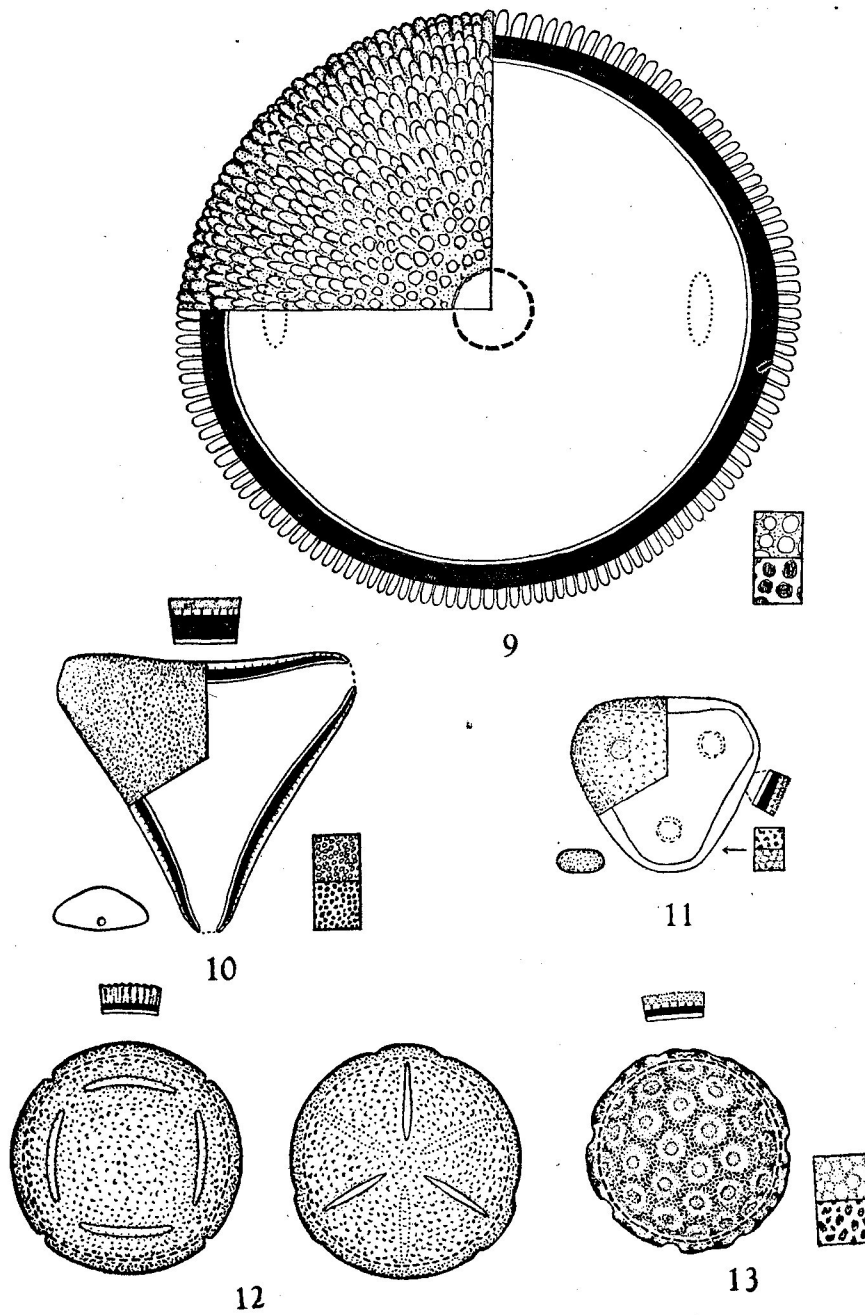
Darlingia spectatissima (fig. 10; Proteaceae): grains subisopolar, zonoporate (3), oblatoid [$22 (17+5) \times 38 \mu$], triangular in polar view. Pores tenuimarginate, their diameter about 5μ .

Exine about 3.3μ thick. Sexine thinner than nexine, probably tectate. Tegillum slightly thicker than length of supporting bacula.

N.B. "Oblatoid" grains (Erdtman l.c., p. 465) are those subisopolar grains which would be oblate if they were isopolar. A size expressed by the figures $22 \times 38 \mu$, refers to a normal (isopolar) oblate grain, but if expressed by the figures $22 (17+5) \times 38 \mu$, to a subisopolar, biconvex (cf. above under *Capsicodendron*) grain.



Text-fig. 2—*Peumus boldus*. 3—*Hedycarya arborea*. 4—*Capsicodendron pimenteira*. 5—*Lemna gibba*.
6—*Oxalis acetosella*. 7—*Myriophyllum spicatum*. 8—*Cordia alliodora*.



Text-fig. 9-13—*Franklandia fucifolia*. 10—*Darlingia spectatissima*. 11—*Anacolsa lutea*.
12—*Drymaria cordata*. 13—*Salsola tragus*.

Anacolosa lutea (fig. 11 ; Olacaceae ; Fiji, Smith 914) : grains 2-zonoporate, oblate ($14.5 \times 23 \mu$), rounded triangular in polar view. Pores six (opposite the rounded corners and arranged in two zones, with three pores in each), circular (diameter about 3μ). Diameter of polar apoporia about 10μ . Height of equatorial apoporium (*vide postea*) about 11.5μ .

Exine at equator about 2.5μ thick except at the rounded corners where the thickness is only about 1μ . Equatorial sexine slightly thinner than nexine (at least where the exine attains its maximum thickness).

N.B. The grains in *Anacolosa lutea* have previously been referred to as 6-forate or 3-diploforate (Erdtman 1952, p. 294). *Dumasia truncata* (Ikuse 1956, p. 96 and pl. 20, fig. 2) also has 2-zonoporate grains. The term apoporium has been defined (Erdtman *l. c.*, p. 460) as a polar area delimited towards the equator by the polar limits of the mesoporia. If, however, there are two (or several) poriferous zones the term apoporium may, as above, be used in a broader sense.

Drymaria cordata (fig. 12 ; Caryophyllaceae ; cult. Uppsala, Sweden) : grains pancolpate (12-colpate), diameter 32μ . Colpi about $13 \times 1.5 \mu$. Apices of adjacent colpi about 5μ apart.

Exine about 2.5μ thick, crassisexinous.

Salsola tragus (fig. 13 ; Chenopodiaceae ; cult. Våsterås, Sweden) : grains pantoporate, diameter 27μ . Pores circular (diameter about 1.5μ).

Exine about 2μ thick or slightly less. Sexine about as thick as nexine. Endosexine baculate.

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ADDENDUM

Since the above was written, further suggestions have been made as follows : the monolete, trilete, and cataporate spores in fig. 1, column 3, are monotreme (trema = aperture) and so are also those in columns 4 and 7 ; those in columns 3 are catatrema, those in 4 anatrema, those in 5 zonotreme, and those in 6 pantotreme (pantoporate and pantocolpate respectively).