Taxonomic notes on *Encephalartos ferox* (Cycadales: Zamiaceae), with the description of a new subspecies from Mozambique

PHILIP ROUSSEAU1*, PIETER J. VORSTER2, ABILIO V. AFONSO3 & ABRAHAM E. VAN WYK1

1H.G.W.J. Schweickerdt Herbarium, Department of Plant Science, University of Pretoria, Private Bag X20, 0028 Hatfield, Pretoria, South Africa.
2Department Botany and Zoology, University of Stellenbosch, South Africa, Private Bag X1, 7602 Matieland, Stellenbosch, South Africa.
3Instituto de Investigação Agrária de Moçambique, P.O. Box 3658, 1100 Maputo, Mozambique.

*Author for correspondence. E-mail: pr.philiprousseau@gmail.com

Abstract

Continued work on the systematics of the genus *Encephalartos* has been conducted over a large area along the Mozambican coast to determine the infraspecific variation and taxonomic status of *Encephalartos ferox* populations. The northern extent of the species was also investigated based on anecdotal information that it might represent a new taxon. Several (45) vegetative and reproductive (ca. 40) characters were analysed and used to re-circumscribe *E. ferox*. Literature and iconographic citations are included, as is information on ecology, cytology and conservation as available. No specimens of *E. ferox* were located north of the Save River in the Sofala Province, Mozambique, and the northernmost population of the species was found to be anomalous. Accordingly a new subspecies, *E. ferox* subsp. *emersus*, is recognised based on its invariably emergent stems, shorter unarmed petiole length, shorter leaves, greater number and narrower leaflets (in seedlings as well) which are more closely spaced. Strobili and peduncles in subsp. *emersus* are shorter and narrower with strobili having a propensity towards yellow, while internally the megasporophylls are green, grading to white away from the outer surface. The IUCN Red List status is suggested as Critically Endangered A4d + B1a,b(v) + B2a,b(v), with the threat of illegal over-collection highlighted.

Key words: cycad, Gymnosperm, taxonomy

Introduction

Taxonomy of the African endemic cycad genus *Encephalartos* Lehmann (1834: 1; Zamiaceae, Christenhusz *et al.* 2011), though remaining stable over the past two decades (Haynes 2012, Osborne *et al.* 2012), currently with 65

Accepted by Maarten Christenhusz: 7 Mar. 2015; published: 1 Apr. 2015
Licensed under a Creative Commons Attribution License http://creativecommons.org/licenses/by/3.0
species and two subspecies recognised, has suffered from several taxonomic uncertainties. An over-appreciation at the specific level as espoused by Dyer (1965a) and Vorster (2004), is due to their reluctance to recognise infraspecific ranks, as robust phylogenies—revealing affinities—had yet to be constructed. Even molecular phylogenies remain unsettled (Van der Bank et al. 1998, 2001, Treutlein et al. 2005, Nagalingum 2011), though the latest efforts (Rousseau 2012) have resolved species groupings (2–7 taxa) with adequate support.

FIGURE 1. Known geographical distribution of Encephalartos ferox subsp. emersus (black dot). The thick black line mirrors the coastal range of E. ferox subsp. ferox, with arrows indicating its northern and southern limits.
Hitherto taxonomic work on *Encephalartos* based solely on herbarium material has yielded less than ideal results, as such material lacks characters only discernible in living specimens (e.g. see the critiqued work of Schuster 1932). Field work is difficult because of the scarcity and remote location of species as well as the sporadic nature of reproductive events, which are of diagnostic value. Even in species as relatively widespread (Fig. 1) and common as *Encephalartos ferox* Bertoloni (1851: 264), *in situ* studies have focused predominantly on the more accessible populations at the southern limits of its distribution in South Africa (Dyer 1965a, Dyer & Verdoorn 1966). Even Mozambican botanists have focused their efforts in and around Maputo (ex. Lourenço Marques), rarely venturing as far north as the Inhambane Province where the type is most probably from—as Bertoloni cites many of the other plants collected by Fornasini (the collector and artist for the type) from that area. Little or no formal work has been conducted towards the northern end of the species’ range, with the distribution limits remaining speculative (Capela 2006, Donaldson 2010).

Most of the infraspecific variation exhibited by *E. ferox* is known informally among cycad enthusiasts based on cultivated material of unknown or doubtful origin (Whitelock 2002). Determining the prevalence of character states in an artificial setting is speculative with eventual generalisation on the species rooted in an incomplete understanding of such variation (Vorster 1993). In our continued work on the systematics of the genus, we here re-circumscribe *E. ferox*, and based on our observations, recognise a new subspecies that we name *E. ferox* subsp. *emersus*.

### Study area, materials and methods

Fieldwork in Mozambique was conducted in the Gaza, Inhambane and Sofala provinces where populations were sampled every ca. 100 km along the coastline. Fieldwork could not be conducted in South Africa, as even though permits for the Isimangaliso Wetland Park were secured, the provincial authorities refused permission (though casual observations were made in habitat). All South African specimens cited are thus field accessioned cultivated plants. In total six distinct regions were sampled; individuals $n=59$. At each locality two to three individuals representing both sexes were sampled. As the northernmost taxa were anomalous, the sample size was increased to $n=41$ ($n=19$ reproductive) and the nearest other population to $n=7$. Each individual was analysed for 45 vegetative and, if reproducing, an additional ca. 40 fertile characters depending on sex and seed availability. Where applicable, nine measurements were made with three each direction: proximally, distally, and medially. In the morphological description of the taxa the arithmetic mean ($\bar{x}$) for diagnostic measurements appears in square brackets.

The area of occurrence for subsp. *emersus* was mapped by GPS. This subspecies is associated with distinct vegetated mounds (raised hillocks) emerging from the coastal plain (Fig. 2). Mapping was done by investigating all marginal mounds until two mounds in line were found devoid of individuals. This was then mapped onto Google Earth Pro where polygons were drawn to calculate the surface area. For area of occupancy all mounds of size more than 100 m² were added. For population size several ($n=10$) mounds of intermittent size were counted for all mature individuals, the mean of which was used and multiplied by the number of mounds >100 m² in the area of occurrence.

The herbaria of LMA, LMU, PRE, and PRU were physically visited, while online access was made to K. The living collections of: the National Botanical Garden of Pretoria, Kirstenbosch and the Lowveld; Manie van der Schijff Botanical Garden (University of Pretoria); and numerous private collections were investigated. The descriptive terminology used follows Melville (1957)—except his sporophyll depth, which is here referred to as sporophyll length, and his sporophyll height, which has been modified to include the sporophyll stalk length; leaflet angles follow Grobbelaar (2002).

### Taxonomic treatment

*Encephalartos ferox* Bertoloni (1851: 264). Type:—MOZAMBIQUE. Inhambane: at or near Inhambane, no date, *ex Fornasini* (K!, painting of holotype). (Figs 3–9)


Habit subterranean (Fig. 3A) to emergent (Fig. 4A); rarely 1 to 4 basal offshoots with Donaldson (1998) stating plants in full sun are more prone to suckering; stems 500–1500 mm long, 200–400(–560) mm in diameter, lacking contractile...
tissue (Stevenson 1980); crown glabrous. Cataphylls similar between reproductive and vegetative structures, acuminate to linear, adaxially glabrous, white turning brown with age, abaxially with grey-brown to golden silky indumentum, (35–40–111(–145) × (4–)10–20–(–25) mm. Leaf base adaxially and abaxially ridged, collar absent, (15–)30–60 × (15–)30–60(–80) mm. Leaves (1060–)1280–2323(–2680) mm long, width distal (50–)105–160(–245) mm, proximal (60–)80–135(–180) mm, median (140–)230–320(–350) mm; rachis adaxially ridged to round in longitudinal section, glabrate green with white indumentum, straight to slightly arching, 25–42(50–60) mm in circumference; petiole with 2–12(–14) spines, unarmed petiole (0–)10–65(125–160) mm long. Leaflets (Fig. 5) broadly rectangular to ovate, colliculate, glossy bright to dark green (rarely brown on emergence), (25–)30–53 pairs, undulating transversely, curving abaxially at margins, pungent with (2–)3–6 distal, (0–)2–4 proximal margin teeth with crowding of 2 or 3 teeth at the apex, without overlap (rarely slightly succuous), length × width × spacing: distal (80–)100–145(–160) × 15–35(–45) × (10–)15–35(–40) mm, proximal (20–)52–70(–100) × (13–)17–30(–38) × (20–)27–48(65–75) mm, median 100–193(–195) × (20–)25–52(59–65) × (15–)25–49(55–84) mm, 16–43(–53) veins; pinna-pinna angle distal ca. 70–180º, proximal ca. (90–)150–180º, median ca. (75–)150–180º, pinna-rachis angle distal ca. 30–45(–75º), proximal ca. 45–75º, median ca. 45(–75º), swing angle distal ca. 0–15º, proximal ca. 0–35º, median ca. 0–15º. Micro- and megastrobili dissimilar. Microstrobili (Figs 3B, 6A, B) fusiform, red to orange or yellow, up to 5 produced successively, elongating from 150–235 mm to 320–500 (single strobilus measured at 960) mm long at maturity, 215–355 mm in circumference; peduncle (160–)200–370(–440) × 80–105(–180) mm; microsperophylls (Fig. 6C) rhomboid 4 to 5 facets, glabrous, length × width × height: distal 8–14 × 12–30 × 20–35(–40) mm, proximal 6–15 × (12–)15–27(–35) × 16–35 mm, median (7–)10–15 × 20–32 × 25–40(–44) mm, central facet flat, distal (3–)4–11(–15) × (5–)6–17 mm, proximal 4–15 × 10–20 mm, median (4–)5–8 × (7–)9–16 mm, 0(–)4) sterile. Megastrobili (Figs 3C, D, 4B, 6C–E) ovoid to cylindrical, red to yellow, solitary (2, very rarely 3) produced simultaneously, (285–)300–510(–575) × (490–)575–705(–740) mm; peduncle (60–)85–135–140(–125) × 215–219 mm; megasporophylls (Fig. 7A–D) angular transversely obovate (rarely perfectly rhombic), 4 to 6 facets, glabrous, verrucose to rarely smooth, length × width × height: distal (13–)15–25(–30) × (20–)30–45(–50) × (26–)35–50(–62) mm, proximal 20–39(–45) × (20–)35–55(–60) × (20–)42–64(–75) mm, median 20–37 × 35–57(–62) × 58–87 mm; central facet flush to slightly sunken, distal 7–15(–17) × 12–20 mm, proximal (10–)15–20 × (12–)17–25 mm, median 8–14 × (10–)15–20 mm, stalk length distal 5–25 mm, proximal 26–38(–45) mm, median 38–57 mm, lateral lobe length distal 5–15(–17) mm, proximal 25–36(–47) mm, median 27–60 mm, sterile portion 70–120 mm long. Ovule sarcotesta white turning red at maturity; seeds 81 to 577 per strobilus (Phelan et al. 1993), cylindrical, 40–50 × 15–20(–32) mm (Dyer 1956a); kernel 9 to 20-grooved, 24–37 × 14–20 mm. Seedling leaves 86–145 mm long, unarmed petiole (53–)80–103 mm, leaflets 3–5 pairs, length × width × spacing: (30–)40–70(–80) × 8–18 × 5–12 mm, with 6–14 veins.

TAXONOMIC NOTES—Encephalartos ferox is an enigmatic species in the genus with previous phylogenetic placement seeing much difference of opinion. In Osborne et al.’s (2001) molecular analysis of nuclear ITS positioned two replicates as unresolved (but monophyletic 92BP) in a clade of southern African species. Their morphological analysis grouped it with E. gracilis Prain (1916: 181) (the only other species with red strobili) with weak support (59BP) (Treutlein et al. 1998, 2001) using isozymes found it early divergent to Eastern Cape (South Africa) species. Treutlein et al.’s (2005) molecular analysis of nuclear ITS positioned two replicates as unresolved but monophyletic 92BP in a clade of southern African species. Their morphological analysis grouped it with E. gracilis Prain (1916: 181) (the only other species with red strobili) with weak support (59BP) (Treutlein et al. 2005), using three plastid and one nuclear region, found numerous samples to group monophyletically with high support (93BP/1.0PP), though the backbone remains unresolved and as such the closest relative remains speculative—possibly an eastern seaboard South African subterranean-stemmed species (see: Rousseau 2012; Rousseau unpublished data).

INFRASPECIFIC VARIATION—Stems of up to two meters (Dyer 1956a) were not encountered and clearly were based on an anomalous specimen. Vegetative states that could not be confirmed from Dyer & Verdoorn (1966) include an unarmed petiole of 240–300(–360) mm, up to five teeth at leaflet apices that seems to be correlated with an increase in width with leaflets of up to 60–80 mm, which is also unconfirmed in situ. The latter character state, often thought diagnostic for the species, seems confined to populations in the southernmost part of its distribution range. Reproductive states of up to 10 to 13 microstrobili (Donaldson 1998) and five or six megastrobili (Whitelock 2002) could not be confirmed and must constitute extreme rarities or a reflection of higher resources availability in cultivation. The prevalence of red strobili has been rejected in the case of males as the majority of microstrobili were orange, while megastrobili have very few exceptions ranging from orange to yellow. However, this should not be over-weighted as these simply represent a cline in pigmentation from red to yellow, although an allelic genetic component cannot be discounted with the current data. Studies to determine whether the colour of individual's strobili can change in situ from red to yellow, although an allelic genetic component cannot be discounted with the current data. However, as detailed fieldwork has not been conducted in the far southern parts of the species’ range (following the expanded dataset and methodology), consistent diagnostic character states may yet be revealed.

Although the southernmost populations have several distinguishing character states, the majority of character states are shared with other populations. Additionally no ecological separation occurs. As such the resurrection of the name E. kosiensis is not justified. However, as detailed fieldwork has not been conducted in the far southern parts of the species’ range (following the expanded dataset and methodology), consistent diagnostic character states may yet be revealed.
FIGURE 2. Encephalartos ferox subsp. emersus: habitat. A. Plants of subsp. emersus (in foreground) growing among the palm, Phoenix reclinata, on a raised mound (ancient termitarium); note surrounding low-lying floodplain with scattered raised mounds covered by woody vegetation. B. Floodplain with circular mounds covered by Phoenix reclinata and other woody plants, including our cycad. The permanently high water table is reflected by ponds (foreground) scattered throughout the area. Photographs: P. Rousseau (A), P.J. Vorster (B).
No plants were located north of the Save River in Mozambique. The closest population to this geographic boundary between the Inhambane and Sofala provinces, is both ecologically and morphologically distinct. Moreover, the closest other *E. ferox* population is ca. 80 km further south, which represents a significant gap in a near-continues range of ca. 1000 km, a discontinuity that is unusual in other members of *Encephalartos*. Hence this population is here recognised as a new subspecies.

Key to the subspecies of *Encephalartos ferox*

1. Stems subterranean (rarely emergent >1 m); median leaflet width 33–52(59–65 rarely up to 80) mm; median leaflet length × width ratio 3.3:1.0; strobili predominantly red to orange; megasporophylls concolourous externally and internally; red to rarely orange or yellow; seeding leaflet width 10–18 mm; ranging from northeastern KwaZulu-Natal (South Africa) to south of Vilanculos (Mozambique), associated with FZ Vegetation type 20: Miombo Woodlands on Lake Basin, Sul do Save Sands, Vegetation type 14b: Littoral dunes (Wild & Barbosa 1968), and CB 1: Maputaland Coastal Belt (Mucina & Rutherford 2006). .

   - Stems emergent, 1.0–1.5 m long above ground; median leaflet width (20–)25–34 mm; median leaflet length × width ratio 5:1; strobili predominantly yellow; megasporophylls discolourous, externally yellow, internally green grading to white away from the outer surface, rarely concolourous red; seeding leaflet width 8–13 mm; with a restricted range from north of Vilanculos to south of the Save River, associated with FZ Vegetation type 44 (Wild & Barbosa 1968): Deciduous Tree Savanna with Palms (badly drained, lowland), confined to raised mounds (hillocks). .

---

**Encephalartos ferox subsp. ferox** (Figs 3, 5A, B & 7C, D)

**Literature citation (descriptions):**—Prain (1917); Hutchinson & Rattray (1933); Schuster (1932); Ogilvie (1939); Lewis (1960); Dyer (1965a); Dyer & Verdoorn (1966); Giddy (1980); Osborne (1987); Goode (1989); Norstog & Nicholls (1997); Heibloem (1999); Goode (2001); Grobbelaar (2002); Jones (2002); Whitelock (2002); Cooper & Goode (2004).


**Habitat:**—*Encephalartos ferox* subsp. *ferox* is near-endemic to the Maputaland Centre of Endemism (Van Wyk & Smith 2001). It is found in Miombo Woodlands on Lake Basin and Sul do Save Sands (Fig. 3A), *Brachystegia spiciformis* Bentham (1866: 312) (Vegetation type 20, Wild & Barbosa 1968) & Littoral dunes, *Mimusops caffra* Meyer *ex de Candolle* (1844: 203) (Vegetation type 14b, Wild & Barbosa 1968) and CB 1: Maputaland Coastal Belt (Mucina & Rutherford 2006). Rainfall 1000–1250 mm per annum, elevation 20–100 m (Donaldson 2010).

**Distribution:**—SOUTH AFRICA: KwaZulu-Natal; MOZAMBIQUE: Maputo, Gaza, Inhambane. In South Africa it is only found in northeastern KwaZulu-Natal, the main distribution is in the province of Maputo, Gaza and Inhambane in Mozambique. It has not been located north of the Save River despite anecdotal claims by Capela (2006).

**Taxonomic notes:**—See Table 1.

**Phenology:**—Strobili found dehiscent/receptive (day/month): 23/03, 30/04; Inhambane, 29/04; Mapinhane, 29/04; Pomene, 30/4; Paindane, 01/05; Zavora, 02/05; Chipingelele; seed shedding around September.

**Fauna:**—*Porthetes* Schöneherr (1838; 1041) sp. nov. 7 (Oberprieler 1995, Downie *et al.* 2008) is known as probable pollinators, while a novel species of Ertyliadae has been found here at all localities visited. The following Geomitridae (Lepidoptera, Staudte 2001, Staudte & Sihvonen 2014) are also known to feed on the subspecies: *Paraptychodes* Warren (1894: 379) sp., *Diptychis meraca* Prout (1928: 19), *Zerenopsis mo* Staudte & Sihvonen (2014: 27), and *Z. lepida* Walker (1854: 571). Seed sarcotesta is routinely removed (consumed) though the agent(s) responsible are unknown. A
seed dispersal agent seems absent as dispersal is very poor with many crowns littered with seed and even germinating seedlings.

**Etymology:**—Specific epithet from the Latin *ferox* = fierce, alluding to the large almost lobe-like leaflet teeth which are hard, stiff and sharp.


**Strobilus volatile chemistry:**—Of the 19 *Encephalartos* taxa analysed by Suinyuy et al. (2013), *E. ferox* subsp. *ferox* was unique with a volatile composition dominated by alkanes (99%).

**Hybrids:**—No natural hybrids are known as no other species of *Encephalartos* occur within the distribution range of *E. ferox*. Artificial hybridisation between *E. ferox* and other members of the genus—which generally hybridise easily with each other—has been mostly unsuccessful (Vorster, unpublished data). Known artificial hybrids (Vorster, unpublished data) have produced very low yields but include, *E. ferox* × *E. woodii* Sander (1908: 257), *E. ferox* × *E. trispinosus* Dyer (1965b: 112), and *E. ferox* × *E. caffer* Thunberg (1775: 284) (see: Holzman 2005).

**Conservation status:**—Threats include removal for local and foreign horticulture, high burning frequency of grassland habitats kills seedlings and habitat is destroyed for coastal development and agriculture. Cousins et al. (2011) found ca. 10% of the *Encephalartos* material traded at the Warwick traditional medicinal market (Durban) to be *cf. E. ferox* where whole individuals are removed.

**FIGURE 4.** *Encephalartos ferox* subsp. *emersus*: female plant from which the herbarium material designated as type (*Rousseau 1175*) has been collected. A. Emergent plant habit; the distinctive trunk is ca. 1.5 m high. B. Megastrobilus; showing macromorphology. Scale bar = 60 mm. Photographs: P. Rousseau.
**IUCN Red List status:**—In 1997: Rare (Walter & Gillett 1998), in 2003: Least Concern (IUCN 2003), in 2012: Near Threatened (Donaldson 2010). The suggested Red List categorisation is still at Near Threatened as the increased stability (at least in the southern parts) of Mozambique has seen a noticeable rise in the tourism in *E. ferox* habitats, which simultaneously destroys habitats and subsequently sources plants from other areas for cultivation.

**TABLE 1.** Comparison of subspecies of *Encephalartos ferox*. Arithmetic means (\(\bar{x}\)) in square brackets; all measurements in millimeters unless otherwise stated.

<table>
<thead>
<tr>
<th>Character</th>
<th><em>E. ferox</em> subsp. <em>ferox</em></th>
<th><em>E. ferox</em> subsp. <em>emersus</em></th>
</tr>
</thead>
<tbody>
<tr>
<td>Habitat</td>
<td>Coastal forest and thicket on littoral dunes; miombo woodland on coastal sand; usually in partial or complete shade</td>
<td>Savanna-grassland with palms on poorly drained riverine floodplain; plants restricted to raised mounds (termittaria) with woody vegetation in a hygrophilous grassland matrix; usually in full sun</td>
</tr>
<tr>
<td>Stem habit</td>
<td>Subterranean to rarely emergent</td>
<td>Emergent</td>
</tr>
<tr>
<td>Emergent stem length</td>
<td>Up to 1 m [300]</td>
<td>Over 1–1.5 m [716]</td>
</tr>
<tr>
<td>Leaf length</td>
<td>Up to 2.6 m [1818]</td>
<td>Up to 2.35 m [1404]</td>
</tr>
<tr>
<td>Petiole length (unarmed section)</td>
<td>(0–)1530–65(75–160) [64]</td>
<td>(0–)10–43(65–125) [23]</td>
</tr>
<tr>
<td>Number of petiole spines</td>
<td>2–4–6</td>
<td>(6–)8–12–14</td>
</tr>
<tr>
<td>Median leaflet width</td>
<td>33–52 [46]</td>
<td>(20–)25–34 [29]</td>
</tr>
<tr>
<td>Leaflet length/width ratio</td>
<td>3.3:1.0</td>
<td>5.1:1.0</td>
</tr>
<tr>
<td>Leaflet length/width/spacing ratio</td>
<td>3.28:0.99: 1.0</td>
<td>5.9:1.1: 1.0</td>
</tr>
<tr>
<td>Number of leaflet pairs</td>
<td>(25–)30–37–45 [34]</td>
<td>33–52 [41]</td>
</tr>
<tr>
<td>Number of veins/leaflet</td>
<td>(27–)35–43–53 [40]</td>
<td>(6–)16–20(30–42) [24]</td>
</tr>
<tr>
<td>Prevalent colour of strobili</td>
<td>Red to orange</td>
<td>Yellow</td>
</tr>
<tr>
<td>Microstrobilus length</td>
<td>320–440(–960) [427]</td>
<td>430–500 [340]</td>
</tr>
<tr>
<td>Microstrobilus peduncle length</td>
<td>(160–)200–370(–440) [236]</td>
<td>218–315 [249]</td>
</tr>
<tr>
<td>Megastrobilus length</td>
<td>(300–)440–510(–575) [418]</td>
<td>(285–)303–430(–480) [360]</td>
</tr>
<tr>
<td>Megastrobilus circumference</td>
<td>(490–)634–705(–740) [626]</td>
<td>(507–)575–675(–729) [609]</td>
</tr>
<tr>
<td>Megastrobilus peduncle length</td>
<td>(65–)85–117(–140) [91]</td>
<td>60–135 [85]</td>
</tr>
<tr>
<td>Megastrobilus peduncle circumference</td>
<td>(141–)157–219 [177]</td>
<td>125–157(–180) [139]</td>
</tr>
<tr>
<td>Megasporophyll external colour</td>
<td>Red</td>
<td>Yellow</td>
</tr>
<tr>
<td>Megasporophyll internal colour</td>
<td>Red</td>
<td>Green, grading to white away from the outer surface</td>
</tr>
<tr>
<td>Megaspore median height</td>
<td>62–87 [78]</td>
<td>58–62 [57]</td>
</tr>
</tbody>
</table>

*Encephalartos ferox* subsp. *emersus* Rousseau, Vorster & Van Wyk, *subsp. nov.* (Figs 4, 5C, 6, 7A, 7B, 8 & 9)

Most similar to *E. ferox* subsp. *ferox* but distinguished (see Table 1) by its invariably emergent stems (Fig. 4A). Shorter unarmed section of the petiole (Fig. 7A), shorter leaves, narrower (Fig. 8B) and more closely spaced leaflets (Fig. 5C) present from the seedlings stage (Fig. 9I), with more leaflet pairs per leaf. Strobili and peduncles are smaller with a propensity for being yellow (Fig. 4B, 6A, D, E), while megasporophylls are discoulourous, green internally when yellow externally (Fig. 7A). Vegetatively similar but reproductively dissimilar to *E. hildebrandtii* Braun & Bouché (1874: 18) from northern Tanzania and southern Kenya (ca. 2000 km to the north), which is distinguished vegetatively by arborescent stems (2.5–6 m), longer leaves (2–3 m), relatively narrower leaflets (80–350 × 13–45 mm), which are more dentate with up to nine marginal teeth, leaflets not undulating transversely, and the unarmed petiole shorter (10–70 mm).

**Type:**—MOZAMBIQUE. Inhambane: [precise locality withheld], growing on an annual floodplain close to a river on a raised sand mound, -1 m, megastrobilus 26/04, *Rousseau 1175* (holotype PRE, isotypes K, LMA, FTG, PRU [including liquid-preserved megasprophylls]).

TAXONOMIC NOTES ON *ENCEPHALARTOS FEROX* Phytotaxa 204 (2) © 2015 Magnolia Press • 107


**Distribution and habitat:**—This represents the most northerly distribution of the species, as fieldwork as far north as Baia de Sofala in Mozambique has not yielded any specimens. The subspecies is restricted to a single population consisting of two concentrations of individuals separated by ca. 1 km, although the habitat is homogeneous. It is separated from the nearest population of *E. ferox* subsp. *ferox* to the south by ca. 80 km and *E. hildebrandtii* to the north by ca. 2000 km. *Encephalartos ferox* subsp. *emersus* occurs in Vegetation type 44 (Wild & Barbosa 1968), namely Deciduous Tree Savanna with Palms (badly drained lowland, Fig. 2). This vegetation type is found on poorly drained lowland, sub littoral zones as a result of water flowing from the undulation elevations of old quaternary dunes alternation with calcareous plains. Rainfall ca. 800 mm per annum. Outskirts are tree savanna dominated by *Brachystegia spiciformis*, and *Pterocarpus angolensis* de Candolle (1825: 419) amongst others. Floodplains are dominated by *Phoenix reclinata* Jacquin (1809: 27), *Hyphaene crinita* Gaertner (1790: 13), with pools and swamps interspersed, dominated by *Phragmites communis* Trinius (1822: 134), *Nymphaea nouchali* Burman (1768: 120), and *Cyperus* Linnaeus (1753: 44) species, while grasslands have irregular patches of several Paniceae and Andropogoneae grasses, with infrequent bare areas with *Sarcocornia tegetaria* Steffen, Mucina & Kadereit (2009: 453). *Encephalartos ferox* subsp. *emersus* is restricted in this vegetation type to circular soil mounds (Fig. 2A) originating from giant termitaria (Complex 13 of Barbosa 1952) raised above the floodplain with its permanent ponds (Fig. 2B), with up to 20 individuals per mound. These mounds show clear successional vegetation patterns correlated with mound age and size starting with *Phoenix reclinata* and *Hyphaene crinita*, later including species such as a member of maculate *Aloe* Linnaeus (1753: 319), *Erythrina huneana* Sprengel (1826: 243), and finally large trees such as species of *Euclea* Linnaeus (1774: 747) and some of the adjacent savanna elements (Fig. 2B). Elevation -10–8 m with the soils grey and sandy.

**Literature citation:**—Rousseau & Mann (2012).


**Taxonomic notes:**—See Table 1. In the diagnosis above, *E. ferox* subsp. *emersus* is also compared with *E. hildebrandtii*. This may raise the question whether our new taxon should not rather have been described as an infraspecific taxon of *E. hildebrandtii*. Is it not perhaps of hybrid origin, maybe even an allopolyloid involving *E. ferox* and *E.
Encephalartos hildebrandtii? Subsp. emersus is located about 2000 km south from the nearest known populations of E. hildebrandtii, thus genetic exchange between populations of E. ferox and E. hildebrandtii is highly unlikely, at least in relatively recent evolutionary times. This is supported as molecular evidence place E. ferox and E. hildebrandtii in different lineages (Treutlein et al. 2005, Rousseau 2012). Encephalartos ferox subsp. ferox has proved very difficult to hybridise with other members of the genus (Vorster, unpublished data; though E. hildebrandtii has not been tested); we would suspect subsp. emersus to behave similarly. Hitherto polyploidy has not been reported in cycads (Gorelick & Olson 2011), making an allopolyploid origin for subsp. emersus unlikely. The reasons for making the new taxon a subspecies of E. ferox is due to geographic proximity, and the considerable similarity in leaflet and cone morphology. Moreover, the red cones rarely encountered in subsp. emersus would support a close phylogenetic association with E. ferox and not E. hildebrandtii. We mainly make mention of the vegetative similarity with E. hildebrandtii to aid in ex situ identification where geographic data are absent.


**Phenology:**—Strobili found dehiscent/receptive as early as (day/month): 28/02, 02/03, 13/03 with most strobili still immature, to 12/04 with most of the colony mature, to as late as 27/04 where most of the microstrobili are spent. Seed dehiscence is still to be observed but speculated to be around September based on circumstantial evidence and interviews with local people.
Fauna:—Two species of Coleoptera are associated with strobili during pollen shedding and are consistent with other pollinators of *Encephalartos* namely *Porthetes* sp. and a species of Erotylidae. Relationship with those found on *E. ferox* subsp. *ferox* has yet to be established. Damaged leaflets would also suggest the presence of Lepidoptera known to be associated with the genus. Seed coats are routinely eaten, most probably by birds and small mammals as is the case in the rest of the genus. Dispersal is very poor with many seeds never leaving the parental crown and thick stands germinating under parents.


**Etymology:**—Subspecific epithet derived from the Latin *emersus*, meaning “standing above” or “raised up”, in reference to the emergent stems as well as the vegetated soil mounds raised above the surrounding floodplains on which the plants grow.

**Additional specimens examined (paratypes):**—MOZAMBIQUE. Type locality: Rousseau & Mann 29–32, Rousseau & Mann 34–47, Rousseau & Mann 50–55 (PRU), Rousseau 1130B (PRU), Rousseau 1175–1183 (PRU), Rousseau 1250–1256 (PRU).

**Conservation status:**—Subsp. *emersus* does not occur in a protected area and illegal collecting has increased over a three year survey period with some of the removed material traced back to the nearby town’s tourist lodges. Large mature individuals are removed and seem to perform poorly in cultivation. The population size is estimated at 15 individuals per mound, with 70 mounds in the area of occurrence equaling >1000 mature individuals. Area of occurrence is 0.13 km², area of occupancy is 0.05886 km². As the subspecies occurs in a single locality where any stochastic event can eradicate the entire subspecies, and continued pressure from illegal collection is probable, we suggest the IUCN Red List rank of Critically Endangered A4d + B1a,b(v) + B2a,b(v).
FIGURE 8. Encephalartos ferox subsp. emerus: vegetative morphology. A. Leaf; proximal portion. B. Leaf; median portion. C. Leaf; distal portion. D. Median leaflet; abaxial surface, showing distinct parallel venation. E. Cataphyll; adaxial (left) and abaxial (right) views. Scale bar = 40 mm. Voucher: Rousseau 1175. Artist: Lesley Deysel.

Acknowledgments

We thank George Mann for help and companionship during field work, Sten Culverwell for assisting with local interviews, Marco de Canha for the Portuguese translation of the abstract, Hester Steyn for preparing the distribution map and Lesley Deysel for the line drawings. Our grateful thanks to Hermenegildo Matimele and Camila Sousa at the IIAM, Maputo, Mozambique, for plant collecting permits and logistical assistance. The Cycad Society of South Africa, University of Pretoria and the South African National Research Foundation are thanked for financial support.

References

http://dx.doi.org/10.1111/j.1096-3642.1865.tb00186.x


http://dx.doi.org/10.1017/CBO9781107051324.022


http://dx.doi.org/10.5962/bhl.title.538


http://dx.doi.org/10.5962/bhl.title.45011

http://dx.doi.org/10.5962/bhl.title.37656

http://dx.doi.org/10.5962/bhl.title.549

http://dx.doi.org/10.2307/4114417


http://dx.doi.org/10.2307/4115074


