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TWO NEW PARROTS (PSITTACIFORMES) FROM THE LOWER EOCENE FUR FORMATION OF DENMARK

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Abstract: Two new fossil psittaciform birds from the Lower Eocene ‘Mo Clay’ (Fur Formation) of Denmark (c. 54 Ma) are described. An unnamed specimen is assigned to the extinct avian family of stem-group parrots, Pseudasturidae (genus and species incertae sedis), while a second (Mopsitta tanta gen. et sp. nov.) is the largest fossil parrot yet known. Both specimens are the first fossil records of these birds from Denmark. Although the phylogenetic position of Mopsitta is unclear (it is classified as family incertae sedis), this form is phylogenetically closer to Recent Psittacidae than to other known Palaeogene psittaciforms and may, therefore, represent the oldest known crown-group parrot.

Key words: Aves, Psittaciformes, Pseudasturidae, parrots, Palaeogene, Eocene, Denmark.

Parrots (Psittaciformes Wagler, 1827) are very familiar birds to us today, being found in most tropical parts of the world. Even though most extant taxa are found in the Southern Hemisphere (Australasia and South America), parrots as a whole range from India to South-East Asia, sub-Saharan Africa and even to North America [the recently extinct Carolina Parakeet Conuropsis carolinensis Linnaeus, 1758 (Salvadori 1891) and the Thick-billed Parrot of Mexico, Rhynchopsitta pachyrhyncha Swainson, 1827 (Bonaparte 1854)].

No parrots were known from the Palaeogene until about 20 years ago when Harrison (1982) assigned a fossil, Palaeopsittacus georgei, from the Lower Eocene London Clay Formation (UK) to the Psittaciformes. However, Mayr and Daniels (1998) suggested that Palaeopsittacus was anisodactyl or at best facultatively zygodactyl. Dyke and Cooper (2000) later showed it to be made up of unassociated elements and therefore of uncertain affinity. Subsequently, in the light of new material from Messel, Germany, Mayr (2002a) considered the feet of Palaeopsittacus to be preserved in an anisodactyl position, and also remarked on some similarities with Recent Podagaridae (frogmouths), Quercypsitta sudrei Mouter-Chauviré, 1992 and Q. ivani Mouter-Chauviré, 1992 from the Upper Eocene of Quercy, France, were described as parrots and placed in the new family Quercypsittidae by Mouter-Chauviré (1992). Psittacopes lepidus Mayr and Daniels, 1998, comprising two nearly complete skeletons from the Middle Eocene of Messel, was assigned to the Psittaciformes. Dyke and Cooper (2000) described Pulchrapollia gracilis and also assigned it to this order. Finally, a small (15 mm long) cranial bone fragment from the Cretaceous Lance Formation of Wyoming, USA, was argued by Stidham (1998) to be a crown-group psittaciform, although Dyke and Mayr (1999) considered it to be of uncertain taxonomic affinity because of the fragmentary nature of the material and the possibility that it could be from any number of other taxa, such as a caenagnathid-like theropod dinosaur (Dyke and Mayr 1999; see also Waterhouse 2006).

Pseudasturidae Mayr, 1998 (a family of small Palaeogene zygodactyl birds; Mayr 1998) were originally described as being of uncertain taxonomic affinity. Mayr (2002b) reassigned Pulchrapollia gracilis as a pseudasturid (discounting the stout coracoid of the holotype as being from the same taxon as the rest of the material and pointing out similarities between Pulchrapollia and pseudasturid specimens held in a private collection from the same Walton-on-the Naze locality and figured in Mayr 1998 and Mayr and Daniels 1998). In the light of this taxonomic reassignment, he recognised the Pseudasturidae (including Pulchrapollia and a number of other Eocene birds) as stem-group Psittaciformes. This placement for Pulchrapollia corroborated the original analysis of Dyke and Cooper (2000).
Finally, the fossils from the Lower Eocene Fur Formation of Denmark considered in detail herein were briefly mentioned but not formally described and/or positioned taxonomically, as parts of reviews of the ‘Mo Clay’ avifauna by Kristoffersen (2002) and Lindow and Dyke (2006). We describe two new fossil parrots (Psittaciformes) on the basis of this material. One specimen is proposed as a new genus and species of psittaciform, the other as a stem-group psittaciform within the Pseudasturidae.

LOCALITY INFORMATION

As noted above, the two fossils described derive from the Fur Formation (Text-fig. 1) and were collected on the island of Mors by Mr Bent Søe Mikkelsen, the former director of the Moler Museum where the specimens are now housed. Although the Fur Formation is a marine diatomite, classified as a siliceous to clayey siliceous ooze (Pedersen and Surlyk 1983; Pedersen et al. 2004), the fossils come from calcareous concretions within the formation, which formed by precipitation of calcite in pore spaces within the diatomite (Pedersen and Buchardt 1996). Fossil remains within them are preserved in three dimensions because the concretions protected their contents from compaction (Pedersen and Surlyk 1983).

Previously (e.g. Pedersen and Surlyk 1983; Kristoffersen 2002) the formation was considered to be of Paleocene or Paleocene–Eocene age. However, it is now regarded as exclusively Early Eocene (Ypresian) because the Paleocene–Eocene boundary has been identified in the underlying Stolleklint Clay Member of the Ølst Formation (Heilmann-Clausen and Schmitz 2000; Beyer et al. 2001; Schmitz et al. 2004). This is supported by 39Ar/40Ar dating of two volcanic ash layers within the Fur Formation, which yielded ages of 54.5 and 54.0 Ma, respectively (Chambers et al. 2003).

The Fur Formation is temporally equivalent to the two uppermost members of the Ølst Formation (Beyer et al., 2001; Text-fig. 1). On a regional scale, it passes laterally into the clays of the Ølst Formation towards the south and east (Pedersen and Surlyk 1983). Towards the west, it correlates with the offshore Sele and Balder formations of the North Sea Basin (Heilmann-Clausen et al. 1985; Pedersen et al. 2004).

EXPLANATION OF PLATE 1

Figs 1–4. Mopsitta tanta gen. et sp. nov.; FU 110/139, holotype from the Lower Eocene Fur Formation, Denmark; digital photographs of right humerus (coated in ammonium chloride to enhance contrast) in 1, caudal, 2, cranial, 3, right lateral and 4, left lateral views.

Figs 5–8. Pseudasturidae Mayr, 1998 incertae sedis, FU 125 from the Lower Eocene Fur Formation, Denmark; digital photographs of left humerus (coated in ammonium chloride to enhance contrast) in 5, caudal, 6, cranial, 7, right lateral and 8, left lateral views. Scale bars represent 10 mm.
WATERHOUSE et al., Mopsitta, Pseudasturidae incertae sedis
MATERIAL AND METHODS

Our anatomical terminology follows Baumel and Witmer (1993), but has been simplified in English with reference to Howard (1929); all measurements were made using dial calipers accurate to 0.1 mm and DMW produced all of the line drawings. The phylogenetic diagnoses of the fossils are based on Waterhouse (2005).

Institutional abbreviations. ANMH, American Museum of Natural History, New York, USA; ZMUC CN, Zoological Museum, University of Copenhagen, Denmark; FU, Moler Museum, Mors, Denmark; SMF, Forschungsinstitut Senckenberg, Frankfurt am Main, Germany.

SYSTEMATIC PALAEONTOLOGY

Class AVES Linnaeus, 1758
Order PSITTACIFORMES Wagler, 1827
Family incertae sedis

Genus MOPSITTA gen. nov.

Derivation of name. After the 'Mo Clay', which is the local designation for the diatomite of the Fur Formation in which the type specimen was discovered, and Psitta, a diminutive of the Latin Psittacus, a parrot or parakeet.

Type species. Mopsitta tanta sp. nov.

Diagnosis. Mopsitta tanta was a relatively large psittaciform bird, similar in size to the extant Cacatua sulphurea Gmelin, 1788 (Yellow-crested Cockatoo; AMNH 2430, 9040). It can be assigned to Psittaciformes on the basis of: a deep capital groove; pneumatic fossa not expanded distally; cranial margin of humeral head markedly expanded distally in its ventral portion; short deltopectoral crest; well-developed flexor process; wide and deep olecranal fossa.

Mopsitta tanta sp. nov.
Plate 1, figs 1–4; Text-figures 2A, 3A

Derivation of name. Latin, tanta, so much, referring to the large size of the holotype specimen when compared with other Eocene Psittaciformes. The type and only specimen was named ‘Mopsitta tanta’ in a PhD thesis (Kristoffersen 2002) that remains unpublished; hence this name, with its orthographically incorrect specific epithet, must be regarded as a nomen nudum.

Holotype. FU 110/139, a single right humerus, slightly crushed (pieces missing from the shaft, head and deltopectoral crest; Pl. 1, figs 1–4).

Type horizon and locality. Fur Formation (Text-fig. 1), island of Mors, Jutland, Denmark.

Diagnosis. As for genus.

Description. FU 110/139 is a relatively large, robust bone (Text-fig. 3A); its shaft is strongly bowed and approximately square in cross section (slightly wider than deep, although this is probably because of slight crushing of the bone). A large, well-developed, domed humeral head is present. Most of the deltopectoral crest has broken off, but it is possible to observe that the crest is projected distally; the proximal edge of this crest is situated distal with respect to the humeral head, forming almost a right angle with the long axis of the bone. A relatively small bicipital surface is present, with a concave curve to the distal edge. The wide, shallow bicipital furrow extends to meet the ligamental furrow. The external tuberosity is rounded and extends into the humeral head without a depression in between. The capital groove is deep, wide and open, separated distally from the pneumatic fossa by a thick ridge connected to the internal tuberosity. The pneumatic foramen is a large, deep, rounded, highly pneumatized, pit. The brachial depression is a very shallow excavation. The ectepicondylar prominence and entepicondyle form a single continuous surface on the lateral margin of the distal humerus. The large, narrow external condyle extends diagonally towards the median of the shaft. The flexor process is well-developed. A small, roughly circular fossa is present laterally, separating the entepicondyle from the entepicondylar prominence. The external tricipital groove is shallower than the internal tricipital groove.

Comparison. Mopsitta tanta is distinguished from other Palaeogene psittaciforms that have preserved humeri (i.e. Psittacopes lepidus Mayr and Daniels 1998) by its large size, more steeply domed humeral head, more concavely curved bicipital crest, sigmoidally curved shaft, more developed internal condyle and entepicondyle, and less prominent entepicondylar prominence (Text-fig. 2B). It differs from other basal Eocene psittaciforms (e.g. the putative pseudasturid ‘Primobucco’ olsoni Feduccia and Martin, 1976; Houde and Olson 1989; Mayr and Daniels 1998) by having a more rounded humeral head, a larger, more rounded bicipital crest, a sigmoidally curved humeral shaft, and a more robust overall shape (Text-fig. 2C). It also differs from Pseudaturidae in having a pneumatic foramen in the pneumatic fossa, poorly developed external tuberosity, a shorter flexor process, and a large external condyle. It differs from extant Psittaciformes (e.g. Eclectus roratus Müller, 1776: Eclectus Parrot; ZMUC CN 10.6.94) by having a more convex and curved distal edge to the bicipital crest, a more sigmoidally curved shaft (although the shaft is rather strongly curved in Cacatua), and a wider external tricipital groove (Kristoffersen 2002) (Text-fig. 2D).
The specimen also displays some similarities to the Middle Eocene Ibis *Rhynchaeites messelensis* (Ciconiiformes: Threskornithidae) (Peters 1983). However, anatomically *Mopsitta tanta* is clearly different from *Threskornithidae*, and thus from *Rhynchaeites*, on the basis of the characters listed in the diagnosis apart from the first and the last. *Mopsitta* can be further distinguished from *Rhynchaeites* by its size (*Rhynchaeites* is considerably smaller), the shape of the capital groove, and because it has a narrower shaft, a more robust median crest and a narrower external tuberosity.

The Eocene bird *Palaeopsittacus georgei* Harrison, 1982, of uncertain taxonomic affinity as noted above (see also Dyke and Cooper 2000), has a smaller humerus than that of *Mopsitta*: the humerus of *Palaeopsittacus* was also described by Mayr (2002a) as being long and slender, unlike the more robust humerus of *Mopsitta*; it also seems to be somewhat less sigmoidally curved.

Comparison. *Pseudasturidae* resembles Recent Galbulae in their humeral morphology but have a shorter deltopectoral crest (Mayr 2002b). The taxon described here is a different from extant psittaciforms, such as *Chalcopsitta atra* Scopoli, 1786 (Black Lory; ZMUC CN 92.206), in being proportionately longer and much more slender, with a shorter sigmoidally bowed shaft (in lateral view); small, round internal condyle.

**Description.** FU 125 is long and slender and has a slightly sigmoidally curved shaft. The shallow ligamental furrow does not extend further laterally than the humeral head. The lateral edge of the deltopectoral crest is broken, but it seems to have a poorly developed external tuberosity. The bicipital crest is relatively small, with a concave curve along its distal edge. The pneumatic foramen is comparatively large and well excavated, with a robust median crest. The capital groove is a relatively small opening on the proximal surface, extending out onto a large, shallow, pneumatic fossa. A slender shaft, circular in cross section with no visible foramen, extends with a slight sigmoidal curve to the distal end of the bone (lateral aspect). Overall, the entire distal end of the humerus is small. A small, but relatively well excavated, triangular internal tricipital groove is also present. The external tricipital groove is smaller in area than the latter but more deeply excavated. An ovoid, shallow, brachial muscle impression is contained within a larger, shallower, brachial depression.

**Comparisons.** *Pseudasturidae* closely resemble Recent Galbulae in their humeral morphology but have a shorter deltopectoral crest (Mayr 2002b). The taxon described differs from extant psittaciforms, such as *Chalcopsitta atra* Scopoli, 1786 (Black Lory; ZMUC CN 92.206), in being proportionately longer and much more slender, with a smaller deltopectoral crest, a smaller distal end to the humerus with a longer flexor process, and smaller external condyle (Text-fig. 4D).
DISCUSSION

Mopsitta tanta is in many respects, more similar to Recent Psittaciformes than to any other Palaeogene psittaciform. Although it is not absolutely certain on the basis of preserved features (humeral morphology cannot be entirely diagnostic at this level), it is highly likely that Mopsitta tanta is a member of Psittacidae, therefore providing further support to the hypothesis of an early Eocene (or earlier) radiation of Psittaciformes; it is likely that representatives of crown-group Psittaciformes such as Mopsitta, existed in the Early Eocene alongside their stem-group counterparts Pseudasturidae and Quercycsittacidae.

The remains reported here are the most northerly parrot fossils yet known, as well as the first fossil remains from this clade to be found in Scandinavia. Their discovery (along with other Early Eocene psittaciform remains from Europe) suggests an Early Eocene (or earlier) Old World centre of origin for parrots and their relatives because, so far, no Gondwanan Psittaciformes have been found in sediments younger than Middle Miocene (Boles 1993). However, because early parrot fossils from the New World and Southern Hemisphere have not been

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**TEXT-FIG. 3.** A–B, measurements of *Mopsitta tanta* gen. et. sp. nov (FU 110/139) and *Pseudasturidae incertae sedis* Mayr, 1998 (FU 125) respectively, with line diagrams to show where these were taken.

found, a Gondwanan origin cannot be entirely ruled out (see Cracraft 1973, 2001; but also Mayr and Daniels 1998). The timing of the radiation of modern bird groups (Neornithes) remains controversial in evolutionary biology (Dyke and van Tuinen 2004). Many molecular-based studies suggest a Cretaceous origin for modern birds (e.g. Cooper and Penny 1997; van Tuinen and Hedges 2001; Paton et al. 2002), whereas fossil discoveries along with some recent molecular studies seem to suggest a post K/T divergence (e.g. Wyles et al. 1983; Olson 1985; Feduccia 1995, 2003; Ericson et al. 2006). The current restricted molecular data present evidence for a Cretaceous diversification for parrots (Miyaki et al. 1998), but the only way to resolve this problem is to find more fossils (Dyke and Cooper 2000). The presence of basal psittaciforms in Northern Europe, along with other, previously described parrots from the Eocene of Europe (e.g. Mourer-Chauviré 1992; Mayr 1998; Mayr and Daniels 1998; Dyke and Cooper 2000), and no real fossil record of parrots below the K/T boundary (Dyke and Mayr 1999) is certainly consistent with an early Cenozoic radiation for (at least) the psittaciforms (Dyke and Cooper 2000).

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