## Catalogue of American Amphibians and Reptiles.

SALTHE, STANLEY N. 1973. Amphiumidae, Amphiuma.

## Amphiumidae Gray Congo eels

Amphiumidae Gray, 1850:54. Type genus Amphiuma Garden 1773, published by Smith, 1821.

Muraenopses Fitzinger, 1843:34. See comment.

Amphiumidae Cope, 1875:25. Cope regarded himself as the author of this family, disregarding Gray.

Amphiumidae Gray, 1825; Kuhn, 1962:361. See comment.

• CONTENT: One fossil genus, *Proamphiuma*, and one genus, *Amphiuma*, both recent and fossil.

• DEFINITION. The premaxillaries are coossified, premaxillary spines being produced posteriorly dorsally, separating the nasals, ventrally in the roof of the mouth. An "atlas" mandibular ligament passes over the parietal. The occipital condyles are stalked. Vomerine teeth form a row on each side, paralleling the maxillary teeth. The internal choanae are bounded by bone laterally but not posteriorly. The columella is fused to the operculum and is attached to the periotic by a narrow isthmus. The first epibranchial is fused to the first ceratobranchial; the second ceratobranchial is absent, even in hatchlings. The vertebrae are amphicoelous with the centrum somewhat compressed and with the foramen chordae in the upper quarter of the glenoid cavity, which is more ellipsoid than round in outline. There is no ypsiloid apparatus, even though lungs are present.

• DESCRIPTIONS, ILLUSTRATIONS. See Amphiuma. The vertebrae of Proamphiuma are illustrated and described by Estes (1969).

• DISTRIBUTION. At present restricted to the Austroriparian biotic province of Dice (1943) in North America. See fossil record, and also species accounts.

• FOSSIL RECORD. Proamphiuma cretacea, of late Cretaceous age, has been described by Estes (1969) from the Hell Creek Formation in McCone County, Montana. Amphiuma (which see) also has a fossil record, ranging from the late Paleocene to the late Pleistocene.

• PERTINENT LITERATURE. All references are cited elsewhere in this account, or in the genus or species accounts.

• REMARKS. The earliest workers, probably including Linnaeus, thought that Amphiuma was a metamorphic stage in the transformation of Siren into some unknown giant salamander (Rusconi, 1821; Duméril, Bibron, and Duméril, 1854), a notion dispelled by Cuvier (1827). After this it was for a time treated as if it was related to Cryptobranchus because both possess four also noting the similarity to Desmognathus in the "atlas"mandibular ligament and the stalked occipital condyles. After being considered as a completely separate group (Davison, 1895) for some years, Dunn (1922) suggested placing the amphiumids with the salamandrids in the Salamandroidea, mainly on the basis of skull and palatal features, an arrangement followed by subsequent workers (Noble, 1931; Brame, 1967). He also, however (Dunn, 1923), noted that Reed's (1920) study of the otic region formed a basis for allying the amphiumids with plethodontids. Regal (1966) argued that the posterior projection of the vomer reflects a general narrowing of the head and that it may be derived from any larval condition, and proposed removal of the amphiumids from the Salamandroidea, and, on the basis of ear structure and chromosome number (2n = 28) is elsewhere found only in plethodontids), that they be allied to the plethodontids. In that same year, Salthe and Kaplan (1966) found that antigenic determinants on the muscle-type lactate dehydrogenase of Amphiuma are more like those of plethodontids than those of any other group of salamanders.

### COMMENT

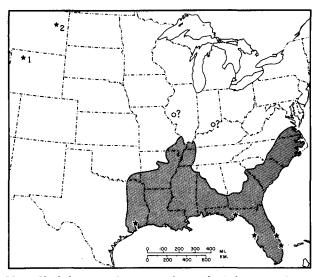
The family name Amphiumidae was first used in 1825 by Gray for a taxon including both Amphiuma and Cryptobranchus. It was not until 1850 (again by Gray) that this name was used for a taxon including only amphiumids, that is, for the taxon as it is now constituted. Therefore, use of Amphiumidae Gray 1825, for example by Kuhn (1962), is incorrect. Tschudi (1838) noted the relationship between Andrias and Cryptobranchus and also that the living representative of the former does not have open gill slits. He therefore broke apart the old family Amphiumidae and gathered the cryptobranchids into a family Tritonides. Apparently being unwilling to erect a monotypic family (there are none in his work of that year), he erected a new family Proteidae, based largely on shared larval characters, to include Amphiuma along with the axolotl, Proteus, Necturus, and the sirenids. It was Fitzinger (1843) who first placed the amphiumids in a family of their own, the Muraenopses, for which he supplied no definition or details of any kind (see comment under Amphiuma). If it were not for this deficiency, his name would under present rules be the family name today. As it was, Gray (1850) took the tack of ejecting the cryptobranchids from his old family, thereby retaining its name.

# Amphiuma Garden Congo eel

- Amphiuma Garden, 1821. ["1773"] p. 599. In Smith, Correspond. of Linn. Type species Amphiuma means Garden, 1773, by monotypy.
- Sireni Linnaeus 1821 ["?1772"] p. 599. In Smith, Correspond. of Linn. Refers to an apparently lost letter from Linnaeus to Garden.
- Chrysodonta Mitchill, 1822:503. Type species Chrysodonta larvaeformis Mitchill, 1822, by monotypy.
- Amphiumina: Bonaparte, 1837:22. Emended name.
- Sirenoidis Fitzinger, 1843:34. Substitute name for part. See comment. Type species Amphiuma tridactylum Cuvier, 1827.

Muraenopsis Fitzinger, 1843:34. Substitute name for part. See comment. Type species Amphiuma didactylum Cuvier, 1827. Amphiuma Linn.: Gray, 1850:162. New combination.

• CONTENT. One fossil species, A. jepseni, and three recent species, A. means, A. tridactylum, and A. pholeter, one or two of which have Pleistocene fossil records.



MAP. Shaded area and questioned extra-limital spots indicate distribution of three Recent species of *Amphiuma*, unnumbered stars mark Pleistocene records for *Amphiuma*. Star 1 marks locality of *Amphiuma jepseni* (late Paleocene); star 2 marks locality of *Proamphiuma cretacea* (Cretaceous).

• DEFINITION. The body is elongated, cylindrical, eel-shaped, with costal grooves. The tail narrows to a point, is com-pressed, and occupies from  $\frac{1}{6}$  to  $\frac{1}{4}$  of the body length. Four ossified limbs are present, very small relative to body size and with few digits, the same number on both fore- and hind-limbs. The head is narrow and more pointed than is usual for urodeles, depressed, and occupying about a tenth of the total length of the body. The eyes are small, without lids, covered with a cuticle continuous with the skin, and are flush with the skin of the head. The lips are thick and extensive, the upper overhanging the lower anteriorly. Ex-ternal gills are absent, but one (vertical) gill cleft remains open on each side; there are four branchial arches. The tongue is flat, indistinct, adherent. Lateral nasal glands are present. Prefrontal and nasal bones are present. The parietals meet in a low sagittal crest, and each has a prominent temporal flange. The vertebrae have prominent muscle crests and two anteriorly-directed hypapophyses (basapophyses) at the anterior ventral extremity of the centrum.

• DESCRIPTIONS, ILLUSTRATIONS. These papers refer to aspects • DESCRIPTIONS, ILLUSTRATIONS. These papers refer to aspects of anatomy that do not differ in the living species. General anatomical studies are Cuvier (1827), Cope (1889), Davison (1895), and Baker (1945). Musculature is described by Davison (1894), Drüner (1904), Maurer (1911), Bruner (1914a), Dunn (1941), Eaton (1936a; 1936b), Auffenberg (1959), Hilton (1959; 1962). The skull and branchial arches are described by Cope (1886), Wiedersheim (1877), Schufeldt (1883), Cope (1886; 1889), and Davison (1895). The nasal (1883), Cope (1886; 1889), and Davison (1895). The nasal region is described by Wilder (1892), Higgins (1919), and Hilton (1951a). The ear region is described by Cope (1888), Norris (1901), Kingsbury and Reed (1909), Reed (1920), Dunn (1922; 1941), Dempster (1930), and Hilton (1950). The teeth are described by Röse (1895), Oltmanns (1952), and With a closed by Röse (1895), Oltmanns (1952), and Hilton (1951b). Glands in the head are described by Wilder (1909), Bruner (1914b), Wilder (1929), and Seifert (1932). The skeleton is described by Cope (1866), Parker (1868), and Wilder (1909), and Seifert (1932). Whipple (1906), while the vertebrae are described by Mivart (1870), Cope (1886), Davison (1897), Hilton (1947b), and Auffenberg and Goin (1959). The lateral line system is described by Norris (1907; 1911), Kingsbury (1896), Escher (1967), Cherge (1902), Cherge (1925), Chezar (1930), and Hilton (1947a). The vascular system is described by Lahti and Churchill (1938), the aortic arches by Baker (1949) and Darnell (1949), the use are are a solution of various respiratory surfaces by Noble (1925), Elkan (1958), and Czopek (1962a), and the heart and cardiovascular dynamics by Johansen (1962; 1964) and Toews (1971a). The lungs have been described by Hilton (1952), Czopek (1962b), Stölk (1962), Willnow (1964), and Tenney and Tenney (1970).

• DISTRIBUTION. See fossil record, and the familial and specific accounts.

• FOSSIL RECORD. Amphiuma jepseni has been described from • FOSSIL RECORD. Ampletational jepsetin has been described from the late Paleocene (Tiffanian) Polecat Bench Formation in Park County, Wyoming (Estes, 1969). This is apparently a somewhat less neotenic form than any of the recent species. For late Pleistocene records see A. means. Amplituma norica from the Pleistocene of Germany (Brunner, 1956) was shown to be spurious by Estes (1969).

• PERTINENT LITERATURE. Spermatogenesis is described by McGregor (1899) and Baker (1962). The former has described the chromosomes, as have Donnelly and Sparrow (1963; 1965) and Dodson (1948). Mirsky and Ris (1957) give the amount of DNA per cell. These papers provide data on cell size, as do Smith (1925), Szarski and Czopek (1965; 1966), and Hartman and Lessler (1964). The thyroid gland and its activity form the subject of papers by Versluys (1925), Baker and Stoudemayer (1951), Kobayashi and Gorbman (1962), Kerkof *et al.* (1963), Moule and Nace (1964), and Prahlad (1970). Hemoglobin kinetics are discussed by Scott (1931), McCutcheon and Hall (1937), and Lenfant and Johansen (1967). Oxvgen consumption is discussed by Smith (1925) the chromosomes, as have Donnelly and Sparrow (1963; 1965) (1967). Oxygen consumption is discussed by Smith (1925) and Morgan et al. (1967), while glycogen utilization in anoxic conditions is studied by Rose et al. (1965) and Rose and Zambernard (1966). The structure and function of the pancreatic islets is studied by Herman and Sato (1970), while the influence of this tissue on glucose metabolism is discussed by Nace and Fucikovsky (1962), Oguri and Nace (1964), and Nace and Blair (1970). Gas tensions in various portions of the blood stream are reported by Toews (1971c). Control of the breathing cycle is the subject of Toews (1971b), while the actual mechanism of obtaining air is discussed by Bruner (1914a, b). Salthe (1965) discusses the ecological and physio-

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• KEY TO THE LIVING SPECIES. Numbers in parentheses indicate the account numbers for the species in this catalogue. 1. Dark ventral coloration .... 2

- Light ventral coloration with dark throat patch; three toes on limbs ..... ......... A. tridactylum (149) 2. Two toes on most limbs; to 890 mm total length .
- . A. means (148) Not more than one toe on limbs; to 305 mm total length \_\_\_\_ . A. pholeter

• REMARKS. See remarks under A. tridactylum for a discussion of the possibility of that taxon being a subspecies of A. means.

• ETYMOLOGY. Garden did not supply an etymological ex-planation for Amphiuma. Duméril, Bibron, and Duméril (1854) noted that other workers were puzzled as to the meaning of this name (evidently Garden was not a strong Greek scholar), and suggested *amphi*, around, and *hyma*, rain, or *amphi*, double, and *hymen*, membrane. The first possibility would presumably refer to the fact that during heavy rains these animals are sometimes caught on roads and other accessible places, the second is obscure and the authors did not elaborate.

#### Comment

Fitzinger's unexplained introduction of two new generic names in 1843 can possibly be explained as a device by which he could escape having a monotypic family. As noted above, Tschudi in 1838 had broken up Gray's old Amphiumidae, and had himself avoided a monotypic family by placing Amphiuma in a family together with several other salamanders that had larval or neotenic traits. The trouble with this suggestion is that Fitzinger does use monotypic families in his classification of 1843, and it also does not take account of his strange treatment of amphiumids in 1826. On page 42 of the Neue Classification he has both Cryptobranchus and Amphiuma as genera making up his family Cryptobranchoidea. On page 66 of that work we find only Cryptobranchus remaining in that family, and Amphiuma has disappeared without comment, to reappear, again without comment, only in the names of the type species of the two new genera in 1843. The name Sirenoidis for the two-toed form did not catch on, but subsequent authors did adopt Muraenopsis for the three-toed form until Ryder (1880) reported specimens with mixed numbers of digits and suggested synonymy of *Muraenopsis* because that name appeared to be based solely on digit number.

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