

Divergent karyotypes of the annual killifish genus *Nothobranchius* (Cyprinodontiformes, Nothobranchiidae)

Eugene Krysanov¹, Tatiana Demidova¹, Bela Nagy²

¹ *Severtsov Institute of Ecology and Evolution, Russian Academy of Sciences, Leninsky prospect, Moscow, 119071 Russia* ² *30, rue du Mont Ussy, 77300 Fontainebleau, France*

Corresponding author: *Eugene Krysanov* (krysanov@sevin.ru)

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Abstract

Karyotypes of two species of the African annual killifish genus *Nothobranchius* Peters, 1868, *N. brienii* Poll, 1938 and *Nothobranchius* sp. from Kasenga (D.R. Congo) are described. Both species displayed diploid chromosome number $2n = 49/50$ for males and females respectively with multiple-sex chromosome system type $X_1X_2Y/X_1X_1X_2X_2$. The karyotypes of studied species are considerably different from those previously reported for the genus *Nothobranchius* and similar to the Actinopterygii conservative karyotype.

Keywords

Africa, chromosome number, karyotype, killifish, *Nothobranchius*

Introduction

Annual killifishes belonging to the genus *Nothobranchius* Peters, 1868 are mainly distributed in eastern Africa but several species are found in central Africa (Wildekamp 2004). They inhabit temporary pools that dry out during the dry season and have specific adaptations for extreme environments. Annual fishes are characterised by specific life history traits of extremely short lifespan and diapause in embryonic development (Furness 2015, Nagy 2015). Their unique biology makes them a model taxon with

which to investigate aging, embryonic development, ecology, and natural selection (Cellerino et al. 2015).

Killifishes of the genus *Nothobranchius* comprise 71 valid species (FishBase 2015). In this genus karyologically were described only 23 species (Arai 2011). These species have variable karyotypes with diploid chromosome numbers ($2n$) ranging from $2n = 16$ for *N. rachovii* Ahl, 1926 to $2n = 43$ for *N. thierryi* (Ahl, 1924) (Scheel 1990). More than 60% of karyotypes in *Nothobranchius* are characterised by a modal diploid number of $2n = 36$ – 38 .

A multiple-sex chromosome system of $X_1X_1X_2X_2/X_1X_2Y$ type has been reported for only one species of *Nothobranchius*, *N. guentheri* (Pfeffer, 1893) with a female karyotype consisting of 36 chromosomes and the male karyotype consisting of 35 chromosomes (Ewulonu et al. 1985).

In this paper, the karyotypes of two species, *Nothobranchius brieni* Poll, 1938 and *Nothobranchius* sp. from Kasenga, were studied, bringing the number of species studied to 25.

Material and methods

Specimens of *N. brieni* were collected from a large ephemeral swamp in the Lualaba drainage, near the village of Bukama in Katanga province (Democratic Republic of Congo, $09^{\circ}11.374'S$ $25^{\circ}51.334'E$) on 2 April 2013 by E. Abwe, B. Katemo Manda, and B. Nagy, whereas specimens of *Nothobranchius* sp. from Kasenga (*Nothobranchius* sp. 'Kasenga') were collected in an ephemeral swamp in the Luapula drainage, near Kasenga, a village in Katanga province (D.R. Congo, $10^{\circ}31.360'S$, $28^{\circ}27.368'E$) on 17 April 2015, by E. Abwe, A. Chocha Manda, B. Katemo Manda, and T. Popp (Fig. 1).

Cytogenetic analysis

Chromosomes were prepared according to the Kligerman and Bloom method (1974). The chromosome preparations were obtained from head kidney tissue. Before preparation fish were treated intraperitoneally with 0.1% colchicine for 3–4 hours. The hypotonisation lasted 20–30 min at room temperature in 0.075 M KCl. Then tissue samples were fixed in 3:1 methanol : acetic acid for 24 hours. Six specimens of *N. brieni* (three males and three females) and three specimens of *Nothobranchius* sp. 'Kasenga' (one male and three females) were karyotyped with this method. Meiotic chromosome preparations of *N. brieni* were acquired from testes by the same technique.

Slides were dried by air and stained with 2% Giemsa solution in phosphate buffer at pH 6.8 for 10 min. Karyotypes were analysed under microscope "AxioImager" Karl Zeiss (Germany) equipped with CCD camera and "KaryoImage" Metasystems Software (Germany). In each specimen the chromosome number and type was determined on metaphase plate. Chromosome morphology was determined according to Levan et



Figure 1. Localities of specimen collections in the Democratic Republic of Congo (1 *Nothobranchius brieni* 2 *Nothobranchius* sp. 'Kasenga').

al. (1964). The chromosomes were classified as metacentric (M), submetacentric (SM), and acrocentric (A). To determine the fundamental number (NF), chromosomes of the M and SM groups were considered bi-armed and those of group A as uni-armed.

Results

The diploid chromosome numbers of *N. brieni* were $2n = 49$ for males and $2n = 50$ for females with NF = 50/50 respectively. The female karyotype consisted of 25 pairs of acrocentric chromosomes gradually decreasing in size (Fig. 2a). The male karyotype consisted of 23 pairs of acrocentric chromosome and one bi-armed pair and two unpaired acrocentric chromosomes (Fig. 2b). In the first meiotic chromosomes during spermatogenesis 23 bivalents and a trivalent were observed at diakinesis (Fig. 2c).

The karyotype *Nothobranchius* sp. 'Kasenga' had diploid number $2n = 49$ for males and $2n = 50$ for females with NF = 68/68 respectively. The female karyotype had two pairs of metacentric, seven pairs of sub-metacentric, and 16 pair of acrocentric chromosomes varying in size from large to small (Fig. 3a). The male karyotype had 23 pair of chromosomes similar to the female with one bi-armed and two unpaired acrocentric chromosomes (Fig. 3b).

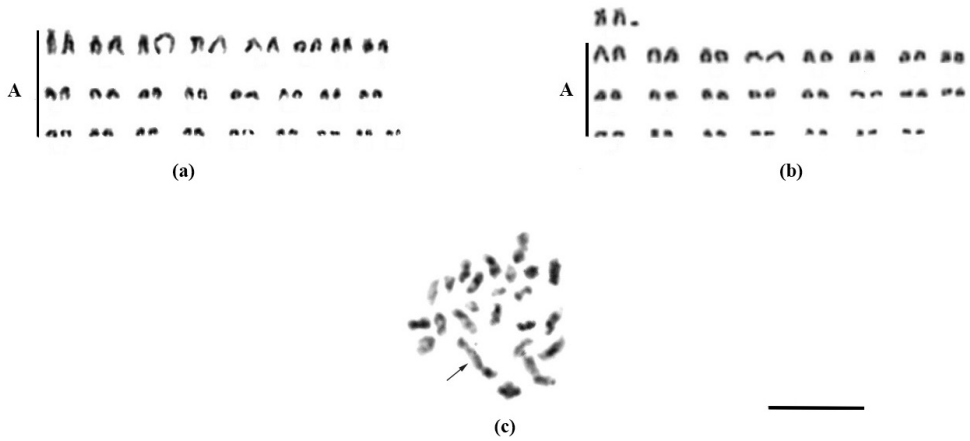


Figure 2. Karyotypes of *Nothobranchius brieni* **a** somatic chromosomes of female **b** somatic chromosomes of male **c** meiotic metaphase I (testicular). (A – acrocentric chromosomes). Note trivalent chromosome (arrowed). Scale bar: 10 μ m.

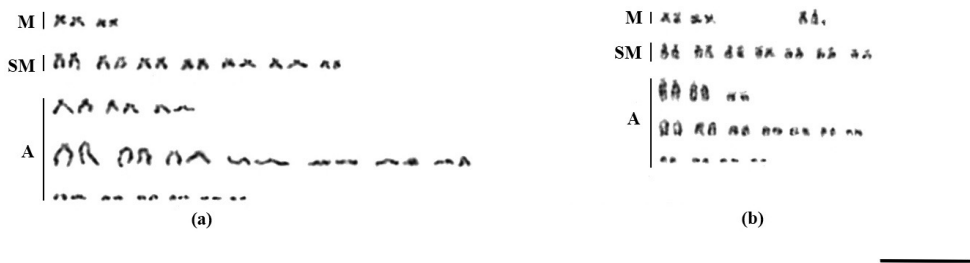


Figure 3. Karyotypes of *Nothobranchius* sp. ‘Kasenga’ **a** somatic chromosomes of female **b** somatic chromosomes of male (M – metacentric chromosomes, SM – submetacentric chromosomes, A – acrocentric chromosomes). Scale bar: 10 μ m.

Discussion

Cytogenetic characteristics

The described karyotypes stand apart from those already reported for species of genus *Nothobranchius*. The karyotype of *N. brieni* has the chromosomal number $2n = 49/50$ and 25 pairs of uni-armed chromosomes in female (50A) and 23 pairs of uni-armed homomorphic and three heteromorphic chromosomes in male (1M + 48A). The karyotype of *Nothobranchius* sp. ‘Kasenga’ has the same diploid number $2n = 49/50$ but a different karyotype structure possessing metacentric, sub-metacentric, and uni-armed chromosomes with 4M + 14SM + 32A for females and 5M + 14SM + 30A for males, while other species of the genus have a considerably lower modal diploid number of only 36 chromosomes (Table 1).

Table 1. The diploid number (2n) of *Nothobranchius* species (from Arai 2011 with modifications). *sex chromosome system of the type $X_1X_2Y/X_1X_1X_2X_2$.

Species	2n	References
<i>N. brienii</i> Poll, 1938*	49♂/50♀	Current study
<i>N. eggersi</i> Seegers, 1982	36	Scheel 1990
<i>N. elongatus</i> Wildekamp, 1982	38	Scheel 1990
<i>N. foerschi</i> Wildekamp & Berkenkamp, 1979	34	Ewulonu et al. 1985, Scheel 1990
<i>N. furzeri</i> Jubb, 1971	38	Reichwald et al. 2009
<i>N. guentheri</i> (Pfeffer, 1893)*	35♂/36♀	Ewulonu et al. 1985, Scheel 1990
<i>N. hengstleri</i> Valdesalici, 2007	38	Wildekamp et al. 2009
<i>N. janpapi</i> Wildekamp, 1977	38	Scheel 1990
<i>N. jubbi</i> Wildekamp & Berkenkamp, 1979	34	Scheel 1990
<i>N. kirki</i> Jubb, 1969	36	Scheel 1990
<i>N. korthausae</i> Meinken, 1973	36	Scheel 1990
<i>N. kryanovi</i> Shidlovskiy, Watters & Wildekamp, 2010	18	Shidlovskiy et al. 2010
<i>N. kuhntae</i> (Ahl, 1926)	38	Scheel 1990
<i>N. lucius</i> Shidlovskiy, Watters & Wildekamp, 2010	36	Wildekamp et al. 2009
<i>N. makondorum</i> Shidlovskiy, Watters & Wildekamp, 2010	38	Wildekamp et al. 2009
<i>N. melanospilus</i> (Pfeffer, 1896)	38	Ewulonu et al. 1985
<i>N. microlepis</i> (Vinciguerra, 1897)	24	Scheel 1990
<i>N. palmqvisti</i> (Lönnerberg, 1907)	36	Ewulonu et al. 1985, Scheel 1990
<i>N. polli</i> Wildekamp, 1978	36	Ewulonu et al. 1985
<i>N. patrizii</i> (Vinciguerra, 1897)	36	Ewulonu et al. 1985
<i>N. pienaar</i> Shidlovskiy, Watters & Wildekamp, 2010	34	Shidlovskiy et al. 2010
<i>N. rachovii</i> Ahl, 1926	16	Ewulonu et al. 1985, Kryanov 1992
<i>N. steinforti</i> Wildekamp, 1977	36	Scheel 1990
<i>N. thieryi</i> (Ahl, 1924)	43	Scheel 1990
<i>Nothobranchius</i> sp. 'Kasenga'*	49♂/50♀	Current study

Sex chromosomes

The reduced diploid numbers and heteromorphic chromosomes in males suggest the occurrence of a multiple-sex chromosome system. A trivalent observation in the first meiotic chromosomes in *N. brienii* and the presence of a bi-armed chromosome exclusively in the male karyotype indicate a multiple-sex chromosome system of the type $X_1X_2Y/X_1X_1X_2X_2$. One bi-armed neo-Y chromosome has most likely resulted from the Robertsonian fusion between the Y chromosome and an autosome, as has been described for other fish species (e.g., Kitano and Peichel 2012). In *N. brienii* and *Nothobranchius* sp. 'Kasenga' the Y chromosome is a large metacentric one, and X_1 and X_2 chromosomes are acrocentric of different sizes. The same-sex chromosome system has been reported only for *N. guentheri* (Ewulonu et al. 1985) among the 23 previously karyotyped species.

Karyotype evolution

In the genus *Nothobranchius* and the related *Aphyosemyon* Mayers, 1924 the evolutionary trend to reduce the total number of chromosomes via acrocentric chromosome fusion was specified (Scheel 1990, Völker et al. 2005). This assumption has been confirmed by the data presented in Table 1. According to this hypothesis, basal taxa have higher chromosome numbers and more acrocentric chromosomes while derived taxa have lower numbers of chromosomes with metacentric chromosomes (Agnese et al. 2006). It is widely accepted that the hypothetical ancestral karyotype of teleostean fishes consisted of $2n = 48-50$ acrocentric chromosomes (Ohno et al. 1969, Nakatani et al. 2007). The two species presented in this study have numbers and a structure of karyotype conservative for Actinopterygii fishes (Mank and Avise 2006, Molina et al. 2014). It is supposed that karyotype of *N. brienii* is similar to that of the hypothetical ancestor of the genus *Nothobranchius*. There is a lack of molecular genetic data on this species, therefore we are not able to consider its phylogenetic position within the clade.

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