Chromosomal analysis of some Egyptian diving beetles (Coleoptera: Dytiscidae)

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ABSTRACT
For the first time, the karyological data on nine African dytiscid beetles is represented. The studied species include: *Hydrovatus cuspidatus* (Kunze), *Hydroporus humilis* klug, *Nebrioporus insignis* (Klug), *Nebrioporus walkeri* (vand der Barnden), *Nebrioporus lanceolatus* (Walker), *Agabus biguttatus* (Olivier), *Colymbetes piceus* Klug, *Eretes sticticus* (Linnaeus) and *Hydaticus decorus* Klug. The data providing useful evidence of chromosomal stability of *Hydrovatus cuspidatus* over a wide geographical area. *Hydroporus humilis* showed 17 pairs of autosomes and sex chromosome. All species of *Nebrioporus* have standard number of chromosome 24 pairs of autosomes plus XX/XO sex chromosomes and very little differences have been shown between the three Egyptian species. For *Agabus biguttatus*, the karyotype of 21 pairs of autosomes plus XX/XO appears normal for genus *Agabus*. For *Colymbetes piceus* and *Hydaticus decorus*, it has been found that diploid number of chromosomes of both species are 20 pairs of autosomes plus XX/X0 sex chromosomes. The world-wide species *Eretes sticticus* has been confirmed to be 2n=43 (21+X0).

KEYWORDS: Coleoptera, Dytiscidae, chromosomes, Egypt

INTRODUCTION
Karyotypes of Coleoptera were first reported in the literature late in the 19th century by Carnoy and early in the 20th century by Henking and Stevens cited in (Makino 1951). During recent years there has been an increase in the studies of cytogenetics of Coleoptera in general and Adephaga in particular. Smith & Virkki (1978) listed all published information on Coleoptera cytogenetics up to that date and Serrano & Yadav (1984) summerized published and unpublished chromosome data for the aquatic Coleoptera up to March 1983. They list data for a total of 426 species, the vast majority of which belong to the Carabidae. Most cytological studies on Adephaga were done mainly on the fauna of Europe (Angus 1983, 1986, 1988, 1989; Angus & Pazos 1991; Shaarawi & Angus 1991a,b; Bilton 1992) and Asia (Yadav & Karamjeet 1981a,b,c; Yadav et al. 1984). In spite of many taxonomic studies on African aquatic beetles there is not any published attempt to explore the karyology of the African fauna.

The sex determining mechanism occurs in different forms in Coleoptera (Smith and Virkki 1978). The majority of organisms rely on chromosomes for sex determination. The system usually comprises a pair of chromosomes called the X and Y chromosomes. Normally females have 2X chromosomes (XX) and males XY.

In many Coleoptera the Y-chromosome is very small and associates with the X during meiosis by means of a nucleolus. At first metaphase the small y-chromosome often has the appearance of a small item being parachuted down, using the X-chromosome as a parachute (John & Lewis 1960), and this arrangement is gradually written XYp or Xyp. In many Adephaga including most Dytiscidae so far studied, there is no y-chromosome. Females have XX, and males have only one X-chromosome, this arrangement being written XO. In some genera, such as *Deronectes* Sharp, the X-chromosome fuses with an autosome to give a neo-
XY system (Nilsson & Angus 1992), with X+ autosome becoming neo-X and the autosome alone the neo-Y.

Recently, Bilton (1992) has recorded the multiple sex chromosome, \(X_1X_2Y\) (male) - \(X_1X_1X_2X_2\) (female) to the three species of genus \textit{Noterus} (family: Noteridae, tribe: Noterini) and similar mechanisms are known in Cicindelidae and 2 species of family: Carabidae. Similarly, the chromosomes of Egyptian \textit{Synchortus imbricatus} (Coleoptera: Noteridae) comprised 12 pairs of autosomes plus sex chromosomes which are \(X_1X_2Y\) (male) and \(X_1X_1X_2X_2\) (female). Unlike \textit{Noterus}, in \textit{S. imbricatus} the Y-chromosome is the largest in the nucleus and is largely heterochromatic (Ahmed et al. 1997).

Multiple sex chromosomes occur when extra chromosomes have been incorporated into the sex determining system (Charlesworth et al. 1987). Such situations can arise from the incorporation of autosomes or via the fragmentation or duplication of the original sex chromosomes themselves (White 1973). Such systems are widespread in the animal kingdom, but show no clear pattern in their phylogenetic distribution (Bilton 1992).

MATERIALS AND METHODS

Table 1 gives details of the material used. Adult beetles were kept in the laboratory in small aquaria (plastic sandwich boxes) until they were used.

**Chromosomes preparation:** Chromosomes were prepared by acetic acid spreading (Angus 1982), mainly from mid-gut and testis tissues. This method was modified for use on adult beetles following Shaarawi & Angus (1989).

For mid-gut chromosomes to be available the beetles should have been feeding for at least about 24 hours. This induces mitosis of mid-gut crypt cells, to replace cells lost in the process of enzyme secretion during digestion (Chapman 1982). Meiotic chromosomes can most readily be observed in the testes of sexually mature male beetles.

**C-banding:** C-banding is often very useful, as it demonstrates constitutive heterochromatin. Slides are kept for 24 hours, then treated for about 7 min. in a saturated solution of barium hydroxide at room temperature (\(20-25^\circ\text{C}\)). They are then rinsed in three changes of distilled water buffered to pH 6.8 with Sorensen before being transferred to double strength Salt Sodium Citrate (2X SSC: 0.3 M sodium chloride with 0.003 M trisodium citrate) at 60°C, and left for 1 hour. They are then rinsed in three changes of distilled water at pH 6.8. They are then stained.

**Staining:** All slides, both unbanded and C-banded, were stained in about 1% Giemsa stain in distilled water at pH 6.8. Slides were stained for about 10 min., then rinsed in distilled water (un-buffered), shaken dry and dried vertically in racks.

**Examination and photography:** Preparations were examined using a Leitz Orthoplan Photomicroscope, using a Leitz Fluortar 100/1.32 oil immersion lens and a Zeiss precision interference filter No. 467808 to give a monochromatic green light. Partial closure of the condenser sometimes helps to give a bolder image. Photographs were taken using Agfa-Gevaert Copex Pan film. For most purposes the automatic exposure was set at 9 DIN, but this was increased to 12 DIN for C-banded preparations. The film was developed for 2 min. at 20°C using Agfa-Gevaert G 141 C developer, mixed 1 part developer to 4 parts water. Prints were made at a magnification of 3000 x, with No. 3 grade paper suitable for most purposes. C-banded preparations were on softer paper (No.1 or No. 2 grade).
Table 1. Species of dytiscid beetles used for chromosomal analysis

<table>
<thead>
<tr>
<th>Species</th>
<th>Locality of Origin</th>
<th>Tissue used</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hydrovatus cuspidatus (Kunze)</td>
<td>Egypt: 10th of Ramadan, Sinai, St. Katherine.</td>
<td>Mid-gut, Testis, Ovary</td>
</tr>
<tr>
<td></td>
<td>Corsica: Pinarelli near Porto vecchio.</td>
<td></td>
</tr>
<tr>
<td>Hydroporus humilis klug</td>
<td>Egypt: Sinai, St. Katherine.</td>
<td>Mid-gut, Testis, Ovary</td>
</tr>
<tr>
<td>Nebrioporus insignis (Klug)</td>
<td>Egypt: Sinai, St. Katherine.</td>
<td>Mid-gut Testis</td>
</tr>
<tr>
<td>Nebrioporus walkeri (vand der Barnden)</td>
<td>Egypt: Sinai, St. Katherine.</td>
<td>Mid-gut, Testis</td>
</tr>
<tr>
<td>Nebrioporus lanceolatus (Walker)</td>
<td>Egypt: Sinai, St. Katherine.</td>
<td>Mid-gut, Testis</td>
</tr>
<tr>
<td>Agabus biguttatus (Olivier)</td>
<td>Egypt: 10th of Ramadan, Sinai, St. Katherine.</td>
<td>Mid-gut, Testis</td>
</tr>
<tr>
<td>Colymbetes piceus Klug</td>
<td>Egypt: Sinai, St. Katherine.</td>
<td>Mid-gut</td>
</tr>
<tr>
<td>Eretes sticticus (Linnaeus)</td>
<td>Egypt: 10th of Ramadan.</td>
<td>Mid-gut, Testis</td>
</tr>
<tr>
<td>Hydaticus decorus Klug</td>
<td>Egypt: Sinai, St. Katherine.</td>
<td>Mid-gut, Testis</td>
</tr>
</tbody>
</table>

RESULTS

This is the first karyological data on some species of African dytiscid beetles including: Hydrovatus cuspidatus, Hydroporus humilis, Nebrioporus insignis, Nebrioporus walkeri, Nebrioporus lanceolatus (Hydroporinae), Agabus biguttatus, Colymbetes piceus (Colymbetinae), Eretes sticticus, Hydaticus decorus (Dytiscinae). The data are presented in Table 2 and Figures 1-4.

Hydrovatus cuspidatus (Fig. 1): Mitotic preparations are shown in Figs. 1.1-1.4. Corsican material is shown in Figs. 1.1, 2, 3, while Fig. 1.4 represents Egyptian material. The karyotype is identical in both localities. The diploid number of autosomes is 16 pairs and the sex chromosome system is X0 in male and XX in female. The autosomes 1 to 6 are mostly metacentric while 7 to 16 are slightly submetacentric and show apparently gradual decrease in size. The X chromosome is clearly metacentric.

Hydroporus humilis (Fig. 2): Mitotic preparations are shown in Figs. 2.1, 2.2. The diploid number of chromosomes differentiates into 17 pairs of autosomes and sex chromosomes which are X0 in male and XX in female. This results in a karyotype of 2n = 34 + X0 in males and 2n = 34 + XX in females. The autosomes are mainly metacentric or nearly so. The first two pairs of autosomes are almost equal in length and the largest element in the nucleus. The rest of the autosomes show fairly gradual decrease in size. X chromosome is submetacentric and clearly larger than the last autosomal pair.

Genus: Nebrioporus (Fig. 3): Four species have been confirmed to be in the Egyptian fauna. Three species were available for chromosomal analysis, N. walkeri Van den Branden, 1885; N. lanceolatus (Walker 1871) and N. insignis (Klug 1834).

Mitotic preparations are shown in (Figs. 3.1, 3.2 N. walkeri; 3.3, 3.4 N. lanceolatus & 3.5 N. insignis). In all species the preparations are from mid gut and testis of adult beetles as in Table 1. The three species have the same diploid number of chromosomes which is (2n = 24 + X0) in male and (2n = 24 + XX) in female.

The sequence of relative lengths and centromere portions of the autosomes appears to be virtually identical in all three species. The last three pairs of autosomes (pairs 22, 23 and 24) are tiny, more or less a dot-like appearance. Seven pairs of autosomes are clearly acrocentric (pairs 6, 7, 9, 10 and 14-16 in the sequence of decreasing relative lengths) in the three species, in addition to one pair more (pair 3) in N. insignis. On the other hand, the first two pairs of autosomes are apparently metacentric in the three species. In all species X-chromosome is acrocentric and about the same length as the middle sized autosomes.
Table 2: Chromosome number and sex mechanism of some Egyptian dytiscid beetles. Data in parentheses are inferred from information on the opposite sex.

<table>
<thead>
<tr>
<th>Species</th>
<th>Diploid no. of chromosomes</th>
<th>Sex mechanism</th>
<th>Karyotype</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>male</td>
<td>female</td>
<td>Male</td>
</tr>
<tr>
<td>Hydrovatus cuspidatus</td>
<td>33</td>
<td>34</td>
<td>X0</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hydroorus humilis</td>
<td>35</td>
<td>36</td>
<td>X0</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nebrioporus insignis</td>
<td>49</td>
<td>(50)</td>
<td>X0</td>
</tr>
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<td></td>
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<td></td>
<td></td>
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<tr>
<td>Nebrioporus walkeri</td>
<td>49</td>
<td>(50)</td>
<td>X0</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nebrioporus lanceolatus</td>
<td>49</td>
<td>(50)</td>
<td>X0</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Agabus biguttatus</td>
<td>43</td>
<td>(44)</td>
<td>X0</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Colymbetes piceus</td>
<td>41</td>
<td>(42)</td>
<td>X0</td>
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<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Eretes sticticus</td>
<td>43</td>
<td>(44)</td>
<td>X0</td>
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<tr>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Hydaticus decorus</td>
<td>41</td>
<td>(42)</td>
<td>X0</td>
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</tbody>
</table>

Figure 1: Mitotic chromosomes of Hydrovatus cuspidatus. 1, 2, 3, 4 males, mid-gut nuclei, unbounded. The autosomes 1 to 6 are mostly metacentric while 7 to 16 are slightly submetacentric and show apparently gradual decrease in size. The X chromosome is clearly metacentric. 1, 2, 3 Corsican material, 4, Egyptian material. Karyotype with simple sex chromosome system 2n = 2n = 33 (16 + X0) in males and in 2n = 34 (16 + XX) in females. Scale line represents 5μm.

Figure 2: Mitotic chromosomes of Hydroorus humilis. 1, male, mid-gut nuclei, unbounded. 2, female, ovary, unbounded. The autosomes are mainly metacentric or nearly so. The first two pairs of autosomes are almost equal in length and the largest element in the nucleus. The rest of the autosomes show fairly gradual decrease in size. X chromosome is submetacentric and clearly larger than the last autosomal pair. Karyotype with simple sex chromosome system 2n = 35 (17 + X0) in males and 2n = 36 (17 + XX) in females. Scale line represents 5μm.
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**Agabus biguttatus** (Fig. 4): Mitotic preparations are shown in Figs. 4.5, 4.6. The karyotype comprises 21 pairs of autosomes and sex chromosomes which are X0 in male and XX in female. Two pairs of autosomes (pairs 3 and 4) are clearly submetacentric, while all the others are more or less metacentric. The autosomes 1 to 13 show a slight decrease in size, while the rest group of autosomes is about the same size. The X chromosome is submetacentric.

**Colymbetes piceus** (Fig. 4): Mitotic preparations are shown in Fig. 4.4. The karyotype possesses 20 pairs of autosomes and sex chromosomes which are X0 in male and almost certainly XX in female. The autosomes are small to medium sized, mainly submetacentric or nearly so except for first pair that is metacentric. The X chromosome is about the same length as in first autosome but clearly submetacentric.

**Eretes stictcus** (Fig. 4): Mitotic preparations are shown in Figs. 4.1, 4.2. The diploid number of chromosomes contains 21 pairs of autosomes and sex chromosomes that are X0 in male and XX in female. This results in a karyotype of 2n = 42 + X0 in males and almost certainly 2n = 42 + XX in females. The autosomes are differentiated into 11 pairs of metacentric (pairs 1-3, 5-8, 10, 11, 14, 15), 5 pairs of submetacentric (pairs 4, 9, 12, 13, 16) and 5 pairs of acrocentric (pairs 17-21). The X chromosome is relatively of medium size and distinctly submetacentric.

**Hydaticus decorus** (Fig. 4): Mitotic preparations are shown in Fig. 4.3. The karyotype consists of 20 pairs of autosomes and sex chromosome that are X0 in male and almost certainly XX in female. The diploid number of chromosomes is 2n = 40 + X0 in male and almost certainly 2n = 40 + XX in females. The autosomes are mostly metacentric except for pairs 1, 3-5 and 12, 13 which are submetacentric. They show clearly decrease in length with last pair (20) of very tiny size. The first pair is distinctly of second constriction and second one is acrocentric. X chromosome is submetacentric.

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Figure 3: Mitotic chromosomes of genus Nebrioporus. 1, males, mid-gut nuclei, unbanded, N. walkeri. 2, males, unbanded, N. lanceolatus. 3, testis. 5, males, mid-gut nuclei, unbanded, N. insignis. The sequence of relative lengths and centromere portions of the autosomes appears to be virtually identical in all three species. The last three pairs of autosomes (pairs 22, 23 and 24) are tiny, more or less a dot-like appearance. Seven pairs of autosomes are clearly acrocentric (pairs 6, 7, 9, 10 and 14-16 in the sequence of decreasing relative lengths) in the three species, in addition to one pair more (pair 3) in N. insignis. On the other hand, the first two pairs of autosomes are apparently metacentric in the three species. In all species X-chromosome is acrocentric and about the same length as the middle sized autosomes. Karyotype with simple sex chromosome system 2n = 49 (24 + X0) in males and {2n = 50 (24 + XX) } in females. Scale line represents 5µm.
Figure 4: Mitotic chromosomes. 1,2 Eretes sticticus, males, mid-gut nuclei, unbanded. The autosomes are differentiated into 11 pairs of metacentric (pairs 1-3, 5-8, 10, 11, 14, 15), 5 pairs of submetacentric (pairs 4, 9, 12, 13, 16) and 5 pairs of acrocentric (pairs 17-21). The X chromosome is relatively of medium size and distinctly submetacentric. Karyotype with simple sex chromosome system 2n = 43 (21 + X0) in males and {2n = 44 (21 + XX)} in females. 3, Hydaticus decorus, males, testis nuclei, unbanded. The autosomes are mostly metacentric except for pairs 1, 3-5 and 12, 13 which are submetacentric. They show clearly decrease in length with last pair (20) of very tiny size. The first pair is distinctly of second constriction and second one is acrocentric. The X chromosome is submetacentric. Karyotype with simple sex chromosome system 2n = 41 (20 + X0) in males and {2n = 42 (20 + XX)} in females. 4, Colymbetes piceus, The autosomes are small to medium sized, mainly submetacentric or nearly so except for first pair which is metacentric. The X chromosome is about the same length as in first autosome but clearly submetacentric. Karyotype with simple sex chromosome system 2n = 41 (20 + X0) in males and {2n = 42 (20 + XX)} in females. 5,6 Agabus biguttatus, males, testis nuclei, unbanded. Two pairs of autosomes (pairs 3 and 4) are clearly submetacentric, while all the others are more or less metacentric. The autosomes 1 to 13 show a slight decrease in size, while the rest group of autosomes are about the same size. The X chromosome is submetacentric. Karyotype with simple sex chromosome system 2n = 43 (21 + X0) in males and {2n = 44 (21 + XX)} in females. Scale line represents 5µm.

DISCUSSION

Dytiscids possess simple sex-chromosome mechanisms (Yadav et al. 1984) with predominant formula X0 formula in male (Smith & Virkki 1978) which frequently gives rise to species with neo XY (White 1973). All investigated species have typical formula X0 of sex-chromosome mechanism.

The range of haploid chromosome number in the present investigations shows a wide distribution in subfamily Hydroptorinae: n = 17 in Hydrovatus cuspidatus to n = 25 in the three species, Nebrioporus insignis, Nebrioporus walkeri, and Nebrioporus lanceolatus. Consequently, genus Nebrioporus has the biggest number of chromosomes. Actually, no clear picture with regard to a primitive dytiscid karyotype and the direction it might have taken during the course of evolution emerges from the available data. Smith (1950) has proposed that the primitive coleopteran karyotype contained nine pairs of autosomes. As more karyological information becomes available, it seems that for the Adephaga higher numbers are usual, even in the more primitive groups. More data are needed before any conclusion can be reached.
Hydrovatus cuspidatus: There is no published information on the chromosomes of Hydrovatus. However, Angus (unpublished data) has chromosomes of Corsican material of H. cuspidatus. The Egyptian material agrees with that from Corsica, providing useful evidence of chromosomal stability over a wide geographical area.

Hydroporus humilis: A simple dytiscid karyotype with 17 pairs of autosomes and the XX/XO sex system normal in the Adephaga (Serrano & Yadav 1984).

Balke & Fery (1993) have shown that H. humilis is closely related to H. tessellatus Drapiez, and comparison of the karyotype would be of great interest. Unfortunately, attempts to obtain English H. tessellatus were unsuccessful because of the summer drought of 1995.

There are no published data on Hydrovorus chromosomes, but Angus (unpublished) has 20 pairs of autosomes plus X0/XY for English H. cantabricus Sharp and Swedish H. melananarius Sturm.

Nebrioporus: The chromosome of this genus are of particular interest as it is the one case so far known where a large dytiscid genus appears to have a standard number of chromosomes 24 pairs of autosomes, ranging from medium sized metacentric to small (dot-like) acrocentric plus XX/X0 sex chromosomes (Nilsson & Angus 1992).

The karyotypes of the three Egyptian species Nebrioporus insignis, Nebrioporus walkeri and Nebrioporus lanceolatus confirm to this pattern, and show very little difference between the three species. The fourth Egyptian species, N. cerisyi was not available for study but Nilsson & Angus (1992) mentioned 50 chromosomes in an Israeli female- apparently 24 pairs of autosomes plus XX.

Agabus biguttatus: The karyotype of 21 pairs of autosomes plus XX/XO appears normal for Agabus (Fery & Nilsson 1993 quoting unpublished data from Angus). At present no chromosome data on non-Egyptian Agabus biguttatus are available, but they would clearly be of great interest in view of the ecological peculiarities of some Egyptian Agabus biguttatus.

There are a number of published records of XY sex chromosome systems in Agabus. Thus, Suortti (1971) recorded A. sturmii Gyll. and A. bipustulatus (L.) as having 21 bivalents at meiosis, including Xyp sex-mechanisms; Yadav et al. (1984) recorded Indian A. conspersus Marsham as having 19 pairs of autosomes plus XY, and Smith (1953) recorded A. confinis Gyll. as having 20 pairs plus XY. However, Angus (unpublished) found that English material of the first three species has 21 pairs of autosomes plus XX/X0, and that a Swedish female of A. confinis had 44 chromosomes. There is thus at present no good reason to believe that any Agabus has an Y-chromosome.

Colymbetes piceus: The cytology of the genus Colymbetes Clairville was first studied by Günthert (1910) who described the oogenesis of Colymbetes fuscus Aubé, mentioned 2n = 35-37 (female) and drew comparison between his and Giardina's results (1901). Smith & Virkki (1978) reported C. paykulli Er. as having 17 pairs of autosomes plus Xyp, C. striatus L. as having 19-21 pairs plus XY and recorded polymorphism in C. paykulli. There are no published data on the chromosomes of C. piceus and in this study it has been found that diploid number of chromosomes 2n = 41 (20+X0) (male).

Eretes sticticus: There are two previous records on the chromosome number of Eretes sticticus, one by Joneja (1960) as 2n = 41 (20+X0) (male) and other one by Yadav et al. (1984) as 2n = 43 (21+X0) (male). The chromosome number given by Yadav et al. (1984) agrees with the present study.

Hydaticus decorus: Joneja (1960) recorded the chromosome number of two species of genus Hydaticus Leach: H. fabricii Macl as 2n = 41 (20+X0) (male) and H. leander Rossi as 2n = 45 (22+X0) (male). However, Yadav et al. (1984) reported 2n = 41 (20+X0) (male) for H. leander and gave the chromosome number for two more species, H. viitatus F. as 2n = 45 (22+X0) (male) and H. luczonicus Aubé as 2n = 41 (20+X0) (male). There is no published data...
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on the chromosomes of *H. decorus* and according to this study it has been found a diploid number of chromosomes 2n = 41 (20+ X0) (male).

REFERENCES


Angus RB (1986) Revision of the Palaearctic species of the *Helophorus minutus* group (Coleoptera: Hydrophilidae), with chromosome analysis and hybridization experiments. *Syst. Ent. 11*: 133-163.


الملخص العربي

التحليل الكروموسومي لبعض أنواع خناص الغوص المصرية (فصيلة دايتسبكيداي)

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من خلال هذه الدراسة، تمت تفهيم التركيب الكروموسومي لجميع أنواع من خناص الغوص العالمية والتي تنتشر في البيئة المصرية. وقد جمعت الدراسة أنواع الثلاثة: فيدروم ذاتيات، هيربورس هوميليس، نيربورس السجينس، نيربورس واكاري، نيربورس لانسينطيس، أهجاس بيجاوتانس، كولومبيتيس بيكيروس، وهاديتيس ديكوراس، أرجاس ستنيكاس. وكانت النتائج على النحو التالي:

- تحتوي الأنواع الكروموسومية من نوع ديترودز كاجاسبيتات في 17 زوج من الكروموسومات الجنسية بالإضافة إلى زوج من الكروموسومات الجنسية.
- أن ثلاثة أنواع مصرية من جنس نيربورس تحتوي على 24 زوج بالإضافة إلى الزوج الجنس، وأن هناك اختلافات طفيفة بين أنواع مصرية الثلاثة.
- يحتوي نوع أهجاس بيجاوتانس في 21 زوج بالإضافة إلى الزوج الجنس، وهذا يتفق مع التركيب الكروموسومي لجنس أهجاس بشكل عام.
- يحتوي كل من كولومبيتيس بيكيروس وهاديتيس ديكوراس في 20 زوج بالإضافة إلى الزوج الجنس.

أكتسبت دراسة التحليل الكروموسومي للفصيلة دايتسبكيداي تطبيقاً مع التركيب الكروموسومي لهذا الجنس الواسع الانتشار في العالم (21 زوج جنسي + زوج جنسي).