# Chromosome morphometry of *Camponotus renggeri* Emery, 1894 (Hymenoptera: Formicidae)

Morfometria cromossômica de *Camponotus renggeri* Emery, 1894 (Hymenoptera: Formicidae)

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**Abstract:** Both the amount and the morphology of chromosomes are important aspects for the specific genomic organization of each organism. Data show that chromosomal evolution, which happens in higher rates in eusocial insects, tends to decrease the size of chromosomes during genetic diversification. Ants have a high karyotypic plasticity, and the genus *Camponotus*, one of the most speciose genus among the Formicidae, has little cytogenetic information available regarding its abundance. Therefore, this study aimed to assess morphometrically the karyotype of *Camponotus renggeri*. The chromosomes were obtained from the brain ganglia of prepupae, following a combination of two existent methodologies. Morphometric analysis of the karyotype revealed chromosomes from 0,31 μm to 1,22 μm, which reflects the chromosomal evolution trend towards considerably small chromosomes.

Keywords: Karyotype. Ants. Cytogenetics.

**Resumo:** A quantidade e a morfologia cromossômica são aspectos importantes para organização genômica de cada organismo. Dados mostram que a evolução cromossômica, que ocorre em taxas mais altas em insetos eusociais, tende a reduzir o tamanho dos cromossomos durante a diversificação genética. As formigas apresentam alta plasticidade cariotípica e o gênero *Camponotus*, um dos mais especiosos entre os Formicídeos, apresenta poucos dados citogenéticos em relação à sua abundância. Este estudo teve como objetivo realizar análises morfométricas dos cromossomos de *Camponotus renggeri*. Os cromossomos foram obtidos a partir dos gânglios cerebrais de pré-pupas, seguindo uma combinação de duas metodologias já descritas. A análise do cariótipo revelou cromossomos de 0,31 µm até 1,22 µm e reflete a tendência de evolução cromossômica, em direção a cromossomos consideravelmente pequenos.

Palavras-chave: Cariótipo. Formiga. Citogenética.

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### INTRODUCTION

Chromosomes are units of inheritance organized within the nucleus of eukaryotic cells and their amount and morphology are important aspects for the specific genomic organization of each organism (Cristiano *et al.*, 2013; Cardoso *et al.*, 2014). Both numerical and structural chromosomal variation is important for systematic and evolutionary studies and may lead to species identification and differentiation, favoring phylogenetic updating (Silveira *et al.*, 2006; Barros *et al.*, 2010; Menezes *et al.*, 2014; Correia *et al.*, 2016).

Cytogenetic data shows that chromosomal evolution tends to decrease the size of chromosomes, reducing genetic risks, such as translocation, during genetic diversification (Aguiar *et al.*, 2016; Mariano *et al.*, 2003). The evolution of chromosome in eusocial insects shows higher rates compared to other organisms and some genera may even display intraspecific variations (Lorite & Palomeque, 2010; Ross *et al.*, 2015; Barros *et al.*, 2016).

Ants have a large numerical and structural variety of chromosomes, with species exhibiting 2n = 2 to species with 2n = 120 (Cristiano *et al.*, 2013). The genus *Camponotus*, with 1,042 described species (Bolton, 2020), is one of the most specious genus among Formicidae and it features a considerable karyotypic variety and has little cytogenetic information available regarding its abundance (Aguiar *et al.*, 2017).

Although it is still scarce, morphometric analysis on ants chromosomes is an important tool for the updating of taxonomic status, once it allows the avaluation of measurements and variations of such structures (Gokhman, 2006; Fornel & Estrela, 2012; Cardoso *et al.*, 2017). Therefore, due to ants high karyotypic plasticity, mainly among *Camponotus*, and the lack of morphometric studies, this work aimed to morphometric assess the karyotype of *Camponotus renggeri* (Emery, 1894), that recently has been the subject of a wide discussion (Aguiar *et al.*, 2017), and we believe that our results may be useful as a mean of comparison for understanding chromosomal variations or rearrangements that might be found in further studies.

# MATERIAL AND METHODS

The biological materials were collected nearby Quirinópolis, state of Goiás, Brazil, in four different colonies, found in a preserved fragment of Cerrado surrounded by pasture fields. Chromosomes were obtained from the brain ganglia (BG) of 10 prepupae from each colonie, following a combination of two methodologies (Imai *et al.*, 1988; Guerra & Souza, 2002), not described by other authors so far, which best suits our laboratory conditions.

Using dissecting needles, the BG were removed under a stereomicroscope and transferred to 500  $\mu$ l tubes with colchicine at 0.005% and allowed to rest for one hour to interrupt the spindle fibers. Later, colchicine was removed and replaced by an hypotonic solution, making cells more turgid. After 15 minutes, the hypotonic solution was removed and 10 drops of carnoy fixative (ethyl alcohol, acetic acid, 3: 1) were added as well.

After a 10 minutes rest, the BG were transferred to  $500 \ \mu$ l tubes with fresh carnoy fixative, where they were held for 30 minutes. Then, using dissecting needles, each BG was macerated with a drop of acetic acid at 45% on separate slides and allowed to dry at room temperature for one day and then stained with a solution of Giemsa at 4% for 10 minutes.

The slides were observed using an optical microscope (Leica DM 500  $\circledast$ ) and the eight metaphases with best chromosome scattering were photographed and characterized according to number and morphology, following to the nomenclature proposed by Levan *et al.* (1964). The morphometric analyzes were performed using the ImageJ software, measuring the whole extension of chromosomes, using a 5  $\mu$ m scale, and the resulting data were organized in table.

# **RESULTS AND DISCUSSION**

The chromosome number found for *C. renggeri* was 2n = 40, of which two pairs are submetacentric (SM), 17 pairs are subtelocentric (TS) and one is telocentric (T) (Figure 1). The karyotype corresponds to existent descriptions for the same species, performed in Nova Mutum-MT and Macapá-AP, both in number and chromosomal morphology (Aguiar *et al.*, 2016, 2017), however, no study has provided morphometric data so far.

Morphometric analysis of the karyotype revealed chromosomes from  $0,31 \,\mu$ m to  $1,22 \,\mu$ m (Table 1). The high rate of karyotype evolution in eusocial insects, whose aim is to decrease genetic risks, may explain the reduced size of chromosomes in relation to other insect and animal species (Mariano *et al.*, 2003; Ross *et al.*, 2015; Barros *et al.*, 2016; Aguiar *et al.*, 2016, 2017). It is also noteworthy that individuals of the same species, collected and karyotyped in other regions, may present variations in chromosome morphology, since some organisms may have intraspecific variations (Lorite & Palomeque, 2010; Barros *et al.*, 2016), and therefore might also present variations in size (Cardoso *et al.*, 2018).

Even though there are not many studies about the karyotype of *C. renggeri*, the available informations has shown that it undergoes a considerable rate of chromosome rearrangements, as expected (Aguiar *et al.*, 2016, 2017). Thus, the measurements presented here allow accurate comparisons and might open new paths that'll help further evolutionary investigations.

Chromosome number	size (µm)	Туре			
1	1,22	Т			
2	1,03	SM			
3	0,88	ST			
4	0,86	ST			
5	0,86	ST			
6	0,85	ST			
7	0,84	ST			
8	0,82	ST			
9	0,76	ST			
10	0,73	SM			
11	0,72	ST			
12	0,72	ST			
13	0,71	ST			
14	0,69	ST			
15	0,66	ST			
16	0,59	ST			
17	0,50	ST			
18	0,46	ST			
19	0,42	ST			
20	0,31	ST			

 $5 \,\mu m$ 



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Figure 1. Conventional cytogenetics of mitotic cells of *Camponotus renggeri* (1000X). Bar =  $5 \,\mu$ m.

Table 1. Morphometric analysis of the chromosomes of *Camponotus renggeri*.



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