

Polyandric behavior in *Trichonephila clavipes*

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Introduction

Belonging to the family Araneidae, the species *Trichonephila clavipes* is distributed from the South of the United States to Argentina, and is well adapted to the urban environment (Robinson & Mirick, 1971). The females of these spiders build big, elaborated and symmetrical webs, sometimes reaching one meter in diameter. The sticky silk of their web is known for its great resistance, which causes many insects and even small birds to be captured by it. This abundance of food in their webs does not benefit only the *T. clavipes* females. Smaller males of *T. clavipes* and kleptoparasites, such as spiders of the subfamily Argyroderinae (Theridiidae) (Silveira & Japyassú, 2012), can be seen in their webs.

With strong size sexual dimorphism favoring females (females > males), the *T. clavipes* spiders have a polyandrous mating system (one female with several males), which can be extremely advantageous. Compared to females that copulate with fewer males, polyandry may result in offspring with greater genetic diversity (Hosken, Garner, Tregenza, Wedell, & Ward, 2003). The abundance of males per web can vary, and, although there isn't a clear answer to why we can find more males in a web than in others, in other spider taxa, males exhibit preferences for larger females (Hoeffer, 1994), due to their body condition being related to their ability to carry eggs.

Moreover, *T. clavipes* females are known to group together in clusters of interconnected webs. In spiders, clustering behavior increases foraging efficiency, habitat exploration, location of males, and protection from predators and parasites (Brach, 1977; Buskirk, 1975; Lubin, 1974; Rypstra, 1981).

The females of *T. clavipes* also have a behavior of combining parts of previous meals with their web, and making a string that stays allocated in the center of their web. It is unclear the purpose of this behavior and the ramifications of it, and if its presence or absence affects the females and males of *T. nephila*, and the kleptoparasites present in the web.

Given that, the objective of this research is to understand what factors influence the amount of *T. clavipes* males in one web. Consecutively, my hypothesis are that: (1) The amount of males in a single web is related to the body conditions of the female. (2) The presence of *Argyrodes* repels males of *T. clavipes* and vice-versa.

Material and Methods

Data sampling

The present study was conducted over five days in February 2020, at Fazenda Água Limpa, Brasília - DF, and four days at the Instituto de Biologia (IB). All spiders that were in the main web of the female *Trichonephila clavipes* were collected, as well as the food strings. The following information was collected: spider identification (collectors' initials + spider number), whether the females were in aggregates or not, the quantity of webs (in case of aggregates), time, location, the amount of males *Trichonephila clavipes*, the amount of *Argyrodes* spp., amount of female *Trichonephila clavipes* found in a cluster, web height, presence of a food string, presence of any other tenant in the web not identified at the time, and whether any spiders were feeding at the time of observation. In the laboratory, the food strings were weighted and morphological parameters of all spiders collected in the field were

measured. For females of *T. clavipes*, we measured the width of the cephalothorax, as well as the total length from the beginning of the cephalothorax to the end of the abdomen. For males of *T. clavipes*, and other tenants, only their total length was measured. All arachnids were weighed alive on the same day they were collected.

Data analysis

The dataset generated at the end of the summer course was edited and cleaned in R. The body condition (BC) was calculated as the residuals from the regression between the log of their mass and their total length. An exploratory data analysis was made to understand the overall patterns in our data. I utilized the “chart.Correlation” function from the “PerformanceAnalytics” package to see any possible correlations present in our numerical variables. To check for possible interactions between qualitative variables, several boxplots with variables that could be correlated were plotted.

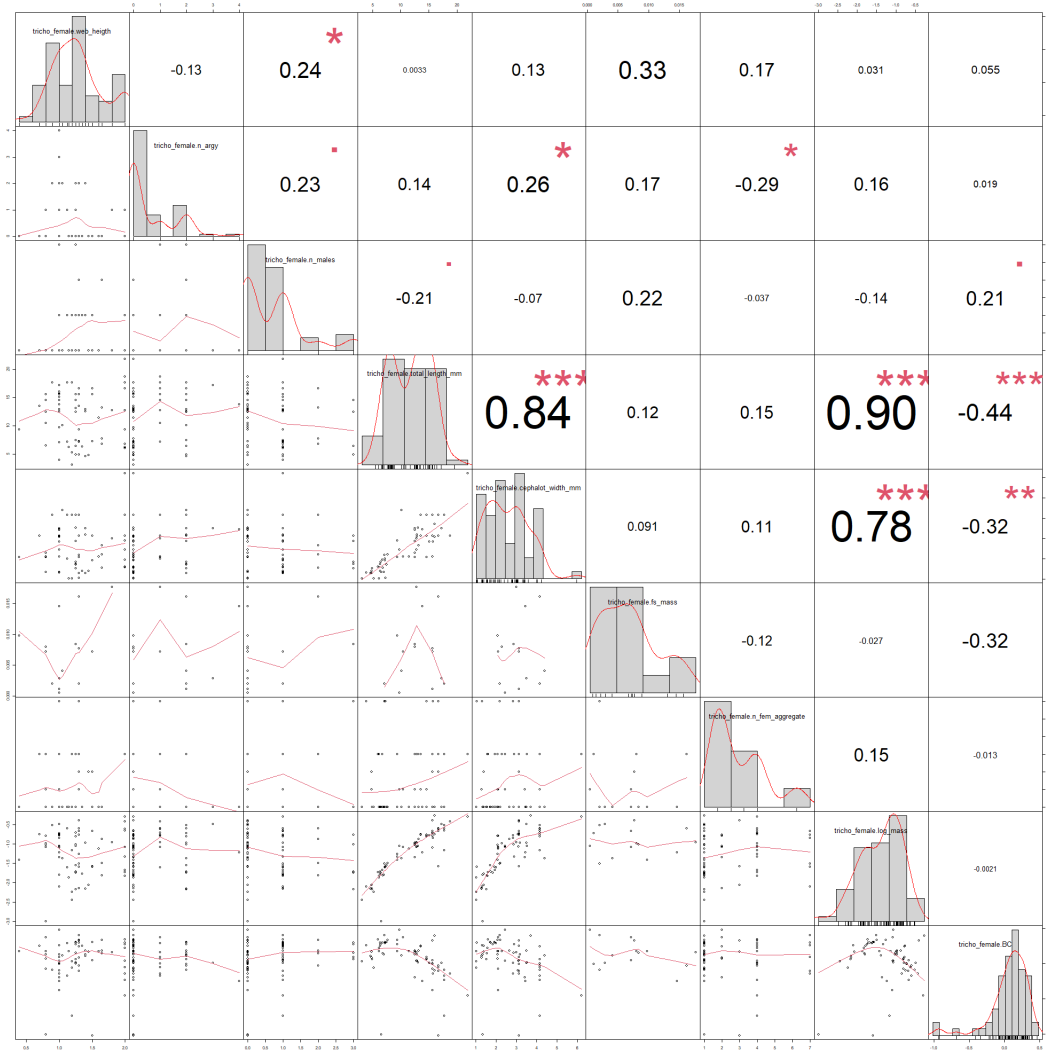


Figure 1: Correlations between the numerical variables found in our dataset. The only meaningful correlations found were between the measurements of the female *T. clavipes*, which are not important to our present study

Subsequently, I used generalized linear models (GLM) to analyze the relationships between the number of males in a web with the *Trichonephila clavipes* females parameters, as well as the spatial parameters (such as the presence of a food string, and if the web is part of an aggregate). Since our response

variable (number of males) has a discrete distribution, I choose the Poisson fit for the GLM. The best model was chosen by the Akaike Information Criterion (AIC), and the selected model was validated, using dispersion and deviation tests, multicollinearity and overdispersion parameters.

Results

Out of 15 models, six showed a delta AIC smaller than 2. Even though I cannot discard models with $\Delta AIC < 2$, I will focus in the two models that had a $\Delta AIC < 1$, tcglm3 and tcglm11. Both of them fitted the assumptions for a GLM.

Table 1: AIC, delta AIC and weights information for each GLM

tcglm	AIC	ΔAIC	df	weight
tcglm3	150.0035	0.0000000	4	0.1771298
tcglm11	150.6450	0.6415723	3	0.1285216
tcglm1	151.6180	1.6145347	5	0.0790132
tcglm7	151.6294	1.6259438	3	0.0785638
tcglm8	151.8719	1.8684128	3	0.0695939
tcglm2	151.8857	1.8822269	4	0.0691148
tcglm14	152.2302	2.2267747	2	0.0581773
tcglm4	152.2498	2.2463034	4	0.0576120
tcglm9	152.4446	2.4411277	3	0.0522646
tcglm5	152.4524	2.4489841	4	0.0520597
tcglm6	152.8994	2.8959295	3	0.0416340
tcglm10	153.0628	3.0593095	3	0.0383682
tcglm13	153.3862	3.3827750	2	0.0326386
tcglm12	153.7285	3.7250048	2	0.0275053
tcglm15	153.7742	3.7707189	2	0.0268838
tcglm16	155.5761	5.5726802	1	0.0109194

The first model (tcglm3) takes in consideration presence of *Argyrodes*, the body condition of the females and if the web that the male is in is part of an aggregate of webs or not. The second model (tcglm11), although it has a higher delta AIC, it has one less parameter (female body condition) than the model with smallest ΔAIC .

Both models present that the number of males increases with the presence of *Argyrodes*, and decreases if the web is part of an aggregate. The model with the smallest ΔAIC also portrays a positive correlation between the number of males and the body condition of the female found in the web.

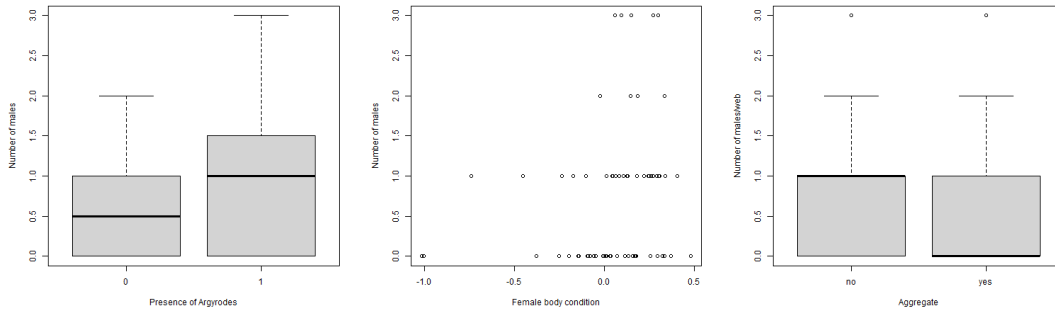


Figure 2: Correlations between the categorical variables present in the selected models and the presence of males. First: Relation between number of males per web and presence of *Argyrodes*. Second: Number of males and female body condition. Third: Number of males and if they are in an aggregate web or a single web.

Discussion

Coinciding with previous studies about *Trichonephila clavipes* (Vincent & Lailvaux, 2006), there seem to be preference by the males for the females with a higher body condition. However, our hypothesis that the males would prefer webs with less *Argyrodes* was proven wrong. Although territorialists between themselves (Christenson & Kenneth C. Goist, 1979), the males not only coexist with other spider species on the female webs, but their numbers increase if there were *Argyrodes* present in the web. We can presume that their territorialism arises from intraespecific competition for the female, and not from interespecific competition for resources. In addition, future analysis could be done to see if the number of *Argyrodes* in a web is also positively correlated to their body sizes.

Although we can see some patterns, we would need posterior statistical analysis to confirm the correlations previously discussed. Moreover, it would be necessary further field work and data sampling, given that the original table doesn't have a big number of observations and it's filled with NAs. However, with this small analysis, we can observe the general patterns regarding the abundance of males *T.clavipes* in a web.

References

- Brach, V. (1977). *Anelosimus studiosus* (Araneae: Theridiidae) and the Evolution of Quasisociality in Theridiid Spiders. *Evolution*, 31(1), 154–161. doi: 10.2307/2407553
- Buskirk, R. E. (1975). Coloniality, Activity Patterns and Feeding in a Tropical Orb-Weaving Spider. *Ecology*, 56(6), 1314–1328. doi: 10.2307/1934699
- Christenson, T. E., & Kenneth C. Goist, Jr. (1979). Costs and Benefits of Male: Male Competition in the Orb Weaving Spider, *Nephila clavipes*. *Behavioral Ecology and Sociobiology*, 5(1), 87–92. Retrieved from <http://www.jstor.org/stable/4599219>
- Hosken, D. J., Garner, T. W. J., Tregenza, T., Wedell, N., & Ward, P. I. (2003). Superior sperm competitors sire higher-quality young. *Proceedings of the Royal Society B: Biological Sciences*, 270(1527), 1933–1938. doi: 10.1098/rspb.2003.2443
- Lubin, Y. D. (1974). Adaptive advantages and the evolution of colony formation in *Cyrtophora* (Araneae: Araneidae). *Zoological Journal of the Linnean Society*, 54(4), 321–339. doi: 10.1111/j.1096-3642.1974.tb00806.x
- Robinson, M., & Mirick, H. (1971). The predatory behavior of the Golden Spider *Nephila clavipes* (Araneae: Araneidae). *Psyche*, 330(3), 701–709.
- Rypstra, A. L. (1981). The Effect of Kleptoparasitism on Prey Consumption and Web Relocation in a Peruvian Population of the Spider *Nephila clavipes*. *Oikos*, 37(2), 179–182. doi: 10.2307/3544463
- Silveira, M. C., & Japyassú, H. F. (2012). Notes on the behavior of the kleptoparasitic spider *Argyrodes elevatus* (Theridiidae, Araneae) 1. *Revista de Etologia*, 11, 56–67. Retrieved from <https://pdfs.semanticscholar.org/3c2f/fd8ca9f5e44df19d89b6fd048dd68af794e9.pdf>
- Vincent, S. E., & Lailvaux, S. P. (2006). Female morphology, web design, and the potential for multiple mating in *Nephila clavipes*: Do fat-bottomed girls make the spider world go round? *Biological Journal of the Linnean Society*, 87(1), 95–102. doi: 10.1111/j.1095-8312.2006.00553.x