

## Research Article

# Mitochondrial PCR-RFLP Assay to Distinguish *Triatoma brasiliensis macromelasoma* from *Triatoma brasiliensis brasiliensis* Subspecies (Hemiptera: Reduviidae)

Daniel Pagotto Vendrami,<sup>1,2</sup> Walter Ceretti-Junior,<sup>1</sup>  
Marcos Takashi Obara,<sup>3</sup> and Mauro Toledo Marrelli<sup>1,2</sup>

<sup>1</sup> Department of Epidemiology, Faculty of Public Health, Universidade de São Paulo, São Paulo, Avendia Dr. Arnaldo 715, 01246-904, SP, Brazil

<sup>2</sup> Institute of Tropical Medicine, Universidade de São Paulo, São Paulo, Brazil

<sup>3</sup> Laboratory of Medical Entomology, Health Surveillance Secretariat, Government Agency for Health, Brasília, DF, Brazil

Correspondence should be addressed to Mauro Toledo Marrelli; [mmarelli@usp.br](mailto:mmarelli@usp.br)

Received 24 June 2013; Accepted 5 November 2013

Academic Editor: Aditya Prasad Dash

Copyright © 2013 Daniel Pagotto Vendrami et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

*Triatoma brasiliensis sensu lato* (s.l.), the main vector of Chagas disease in northeastern Brazil, is a species complex comprising four species, one with two subspecies (*T. brasiliensis brasiliensis*, *T. brasiliensis macromelasoma*, *T. juazeirensis*, *T. sherlocki*, and *T. melanica*), and each taxon displaying distinct ecological requirements. In order to evaluate the genetic relationships among nine *T. brasiliensis* s.l. populations from northeastern Brazil, we analyzed their mitochondrial cytochrome c oxidase subunit 1 sequences and suggested a PCR-RFLP assay to distinguish between *T. b. macromelasoma* and *T. b. brasiliensis* subspecies. All the specimens were morphologically identified as *T. b. brasiliensis*. The resulting phylogenies identified two major clades that are congruent with the geographical populations studied. Based on collection sites and in accordance with type-location, one clade was identified as the subspecies *T. b. macromelasoma*. The second clade grouped *T. b. brasiliensis* populations. Restriction endonuclease sites were observed in the sequences and used in PCR-RFLP assays, producing distinct fingerprints for *T. b. macromelasoma* and *T. b. brasiliensis* populations. The results suggest that these are different species and that gene flow occurs only among *T. b. brasiliensis* populations, possibly associated with human activity in the area.

## 1. Introduction

About 28 million people live in areas at risk of Chagas disease, 11–14.5 million of whom are affected worldwide. *Trypanosoma cruzi*, the pathogen that causes Chagas disease, is found in most South American countries, representing an important cause of heart damage among the economically active population [1]. After a successful chemical control of *Triatoma infestans* (Klugi, 1834), the other main vectors of Chagas causing agent, *Panstrongylus megistus* Burmeister, 1835, *Rhodnius prolixus* Stal, 1859, and *Triatoma brasiliensis sensu lato* Neiva 1911. *T. brasiliensis* kept attracting considerable attention from local entomological surveillance.

*Triatoma brasiliensis sensu lato* (s.l.), found in anthropogenic habitats and considered the main vector in northeast Brazil [2, 3], was recently found to be a species complex that includes *T. b. brasiliensis*, *T. b. macromelasoma* Galvão, 1956, *T. juazeirensis* Costa & Felix, 2007, *T. sherlocki* Papa, Jurberg, Carcavallo, Cerqueira & Barata, 2002, and *T. melanica* Costa et al., 2006. These taxa exhibit wide phenotypic and morphological variability, displaying specific ecological requirements and chromatic patterns [4]. In this respect, accurate species identification is necessary for effective vector control.

The systematic of Triatominae species is based on morphological characters of the adult exoskeleton and male phallic structures [5]. However, insects captured during vector

monitoring and control or received for identification and notification are often immature. Although their characteristics are similar to those of adult individuals, they are difficult to distinguish. The morphology of Triatominae species is not well described; with studies on the immature forms performed for only 40 species, eggs and nymphs described for only 20 species, a key to identify nymphs to the species level has yet to be developed. Available keys are useful and partially applicable to other stages, but specific identification of all live forms remains unresolved [6].

Members of the *T. brasiliensis* complex have been distinguished by analyzing isoenzymes [7], mitochondrial DNA sequences [8], and random amplification of polymorphic DNA-RAPD [9]. In the present study we analyzed the bar-coding *COI* sequences of nine *T. brasiliensis* s.l. populations from northeastern Brazil (States of Pernambuco, Paraíba, and Rio Grande do Norte) in order to identify their genetic relationships. We also conducted a PCR-RFLP assay to distinguish between *T. b. macromelasoma* and *T. b. brasiliensis* subspecies.

## 2. Materials and Methods

**2.1. Sample Collection.** Triatominae were collected by surveillance technicians during active inspections in anthropogenic environments (domestic and peridomestic habitats). Live specimens were collected in nine localities in Northeast Brazil (Table 1, Figure 1) using tweezers, flashlights, and PIRISA. Housed in plastic boxes (7 cm diameter × 8 cm high) lined with folded filter paper, the bugs were transported in coolers to the Culicid and Triatomine Laboratory of the Department of Epidemiology, Faculty of Public Health/USP.

Adults were identified as *T. b. brasiliensis* according to the key by Lent and Wygodzinsky [5]. Nymphs were assumed to be *T. brasiliensis* s.l. because, in addition to the difficulty in distinguishing immature Triatominae based on chromatic and morphological characters, comparative material or a key for nymph identification to species level are not available. After morphological identification, DNA was extracted from individuals from the nine localities and sequenced the 520 bp barcode portion of the *COI* gene (Table 1). Specimens collected in Pernambuco State (even nymphs) were assumed to be *T. b. macromelasoma*, according to Costa et al. [10] and Costa et al. [11].

**2.2. DNA Extraction and *COI* Amplification.** Genomic DNA was extracted from the legs of 10 individual samples of each population using the Qiagen DNeasy Blood and Tissue Kit (Qiagen, Crawley, United Kingdom) following the manufacturer's protocol. The *COI* barcode region was amplified from whole genomic DNA using primers LCOI 1490 (5'-GGT CAA CAA ATC ATA AAG ATA TTG G-3') and HCOI 2198 (5'-TAA ACT TCA GGG TGA CCA AAA AAT-3') [12].

PCR amplification was carried out in a final volume of 50  $\mu$ L containing PCR buffer, 0.2 mM of each dNTP, 2.5 mM MgCl<sub>2</sub>, and 1.25 units of *Taq* polymerase. Initial PCR denaturation was at 94°C for 5 min, followed by 40 cycles of denaturation (1 min) at 94°C, annealing (2 min) at 50°C, and

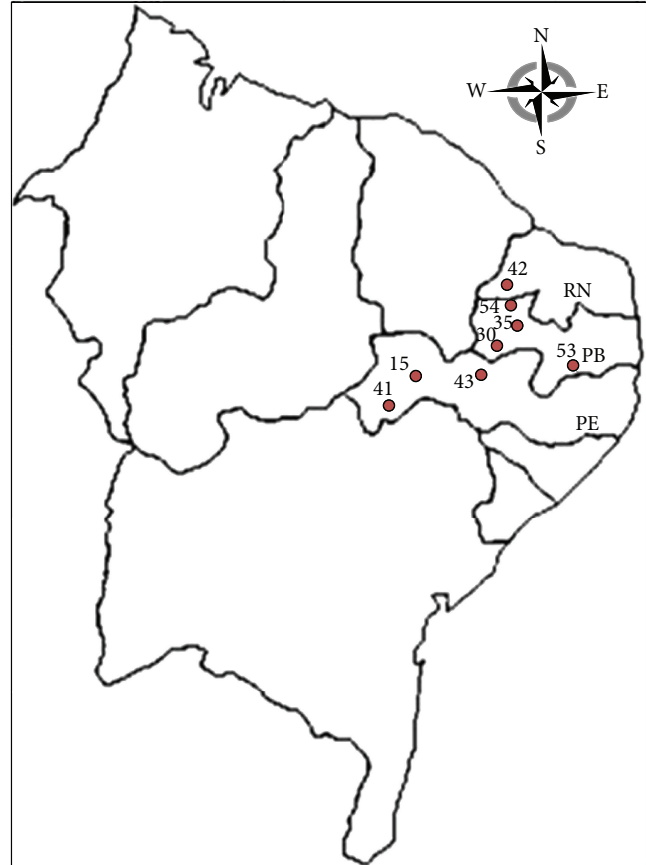


FIGURE 1: Map of northeast region of Brazil indicating the localities of the *Triatoma brasiliensis* populations studied. Paraíba State: Monteiro (30), Mãe d'água (35), Santa Cruz (54), and São José (53). Pernambuco State: Lagoa Grande (41), Salgueiro (15), and Serra Talhada (43). Rio Grande do Norte State: Caicó (42).

extension (2 min) at 72°C. A final extension step at 72°C was performed for 10 min. The amplified DNA was loaded onto 1.5% agarose gel and stained with ethidium bromide. Amplicons were sequenced in both forward and reverse directions using ABI PRISM BigDye Terminator Cycle Sequencing Ready Reaction Kits (Perkin Elmer, Foster City, CA) on an ABI PRISM 3100 Genetic Analyzer/HITACHI.

**2.3. Analysis of DNA Sequence.** Sequences were aligned using Clustal W [13]. One or two representative haplotypes for each population was chosen because of the low variation within populations. Phylogenetic reconstructions were performed by Neighbor Joining and Maximum Likelihood methods (both using the Kimura-3-parameter distance model K81) in MEGA 5.0 [14], and a divergence matrix was constructed under Kimura two parameters (K2P) (Table 2). Maximum Parsimony was carried out using the *branch-and-bound* search option. Phylogenetic analyses included 1000 bootstrap replicates and a *Triatoma sordida* *COI* sequence (Genbank acc. no. AF021213) as outgroup. *Triatoma brasiliensis* s.l. *COI* sequences were also analyzed using NEBcutter version 2.0 [15] to select appropriate endonuclease enzymes.

TABLE 1: *Triatoma brasiliensis* collection sites, coordinates, labels of selected sequences, and GenBank access numbers.

Localities (state)	Coordinates	Label	GenBank
Salgueiro (PE)	08°04'21.60"S 39°07'57.39"W	15 A/B	JQ088297/JQ088298
Monteiro (PB)	07°53'29.40"S 37°07'00.79"W	30 A/B	JQ088299/JQ088300
Mãe d'água (PB)	07°15'10.03"S 37°25'58.25"W	35 A/B	JQ088301/JQ088302
Lagoa Grande (PE)	08°59'07.10"S 40°18'20.46"W	41 A/B	JQ088303/JQ088304
Caicó (RN)	06°27'30.69"S 37°06'09.71"W	42 A/B	JQ088305/JQ088306
Serra Talhada (PE)	07°59'09.46"S 38°17'37.70"W	43 A/B	JQ088307/JQ088308
São José (PB)	06°52'00.00"S 38°38'00.00"W	53 A/B	JQ088309/JQ088310
Santa Cruz (PB)	06°31'33.48"S 38°03'23.52"W	54 A/B	JQ088311/JQ088312
São Francisco (PB)	06°36'53.43"S 38°05'21.89"W	55 A/B	JQ088313/JQ088314

2.4. *Enzyme Restriction Analysis.* Individual *COI* sequences were PCR-amplified using the above parameters and digested in a 10  $\mu$ L reaction with *StyI* (Promega) and *HincII* (New England Biolabs, Ipswich, MA) enzymes. The reaction contained 1  $\mu$ L of 10x buffer, 4  $\mu$ L deionized water, 4  $\mu$ L of amplification product, and 1 unit of restriction enzyme. The digestion mixture was incubated at 37°C for 2 h and then resolved on 2.0% agarose gel.

### 3. Results

Phylogenetic trees derived from the Neighbor Joining, Maximum Likelihood, and Parsimony methods showed similar topologies (data shown for ML tree in Figure 2). Since most of *COI* sequences were identical into and among the populations, the ML tree was constructed using only two samples of each one. This ML tree and the Nucleotide Distance Matrix (Table 2) indicated sequence divergence of up to 4% between the two main clades. The basal clade, with about 4% divergence from the other populations, consisted of the *Salgueiro* population (15A/B). This population was considered to be *T. b. macromelasoma* due to its high sequence divergence compared to the other populations and because it was collected in its type locality [16]. The second clade, with pairwise distances up to 1%, showed that *Pernambuco* populations are more basal, although those from *Serra Talhada* clustered with *Paraíba* populations likely because of the city's proximity to the Pernambuco-Paraíba border. *Paraíba* populations formed a large cluster that also included the *Rio Grande do Norte* population.

Based on sequence analysis, a PCR-RFLP assay was performed to differentiate between the subspecies *T. b. macromelasoma* and *T. b. brasiliensis*. PCR fragment digestion using the *StyI* enzyme produced two restriction fragments (342 bp and 192 bp) in *COI* sequences from *Salgueiro* samples and only one fragment in samples of the other eight populations.

Conversely, the *HincII* enzyme yielded two fragments (297 bp and 240 bp) in all population samples ( $n = 10$  samples of each population) except that from *Salgueiro* (Figure 3). *StyI* and *HincII* enzymes therefore produced distinct fingerprints for *T. b. macromelasoma* and *T. b. brasiliensis*, suggesting that they are different subspecies. The molecular protocols described above are a useful tool in the study of populations and cryptic species, contributing to the identification of insect vectors.

### 4. Discussion

The identification of adult Triatominae based on morphological and chromatic pattern is considered relatively easy for most species; however, this is commonly misguided owing to the wide phenotypic variability within this subfamily. For instance, *Triatoma maculata* Erichson, 1848 and *Triatoma pseudomaculata* Corrêa & Espínola, 1964 which were first treated as members of a same species complex due to morphological similarities [5] thereafter proved to be genetically distant [17–19]. *Panstrongylus herreri* Wygodzinsky, 1948 and *Panstrongylus lignarius* Walker, 1873 in turn, were considered to be separate species until Marcilla et al. [20] and Crossa et al. [21] demonstrated that they are the same species, cytogenetically identical with regard to the second internal transcribed spacer. Another difficulty in identifying genera and species of Triatominae is their extensive chromatic variability. The color of some species such as *Rhodnius* sp. (light brown tones) and *Rhodnius nasutus* Stål, 1859 (pinkish tones) seems to be associated with the color of the palm trees they colonize [22], but others such as *Triatoma rubrovaria* Blanchard, 1843, exhibit well-known 4 chromatic morphotypes [23]. Other studies report the occurrence of natural homoploid hybrids between *T. infestans* and *Triatoma platensis* Neiva, 1913, *T. infestans* and *Triatoma rubrovaria*, and sympatric species of *Phyllosoma* complex and species of the *T. brasiliensis*

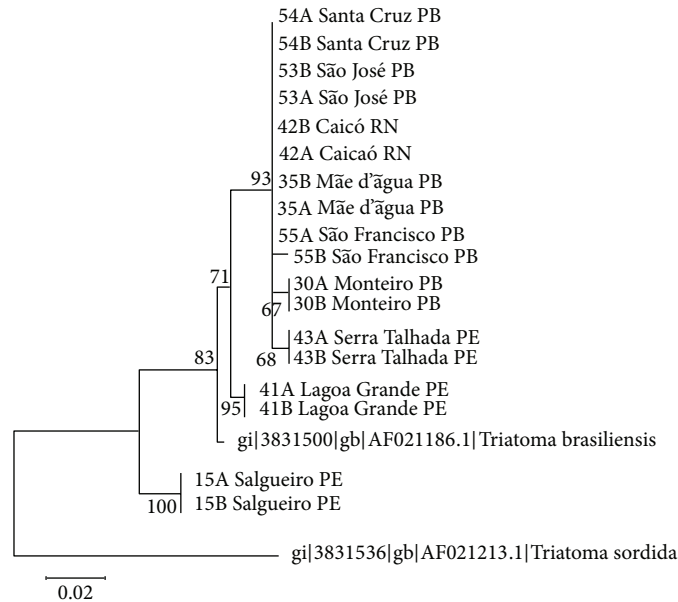


FIGURE 2: Maximum Likelihood phylogeny generated from *COI* sequences using the Kimura-2-parameter model with 1000 bootstrap replicates. Sequences were rooted with *T. sordida* (GenBank acc. no.: AF021213).

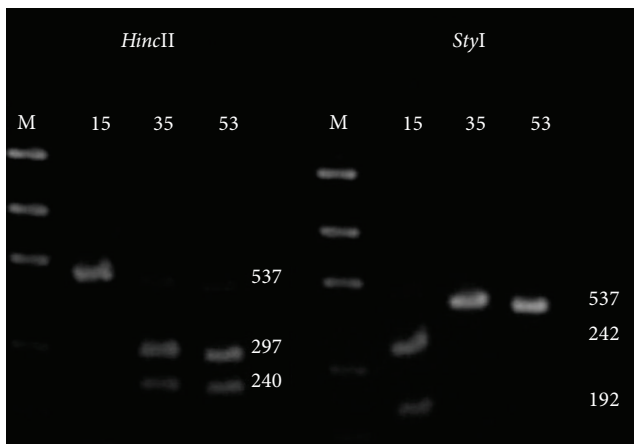


FIGURE 3: PCR-RFLP fingerprint of *T. brasiliensis* from three localities produced with *StyI* and *HincII* enzymes on 2.0% agarose gel. M: DNA ladder 100 bp (New England Biolabs), 15: *Salgueiro*, 35: *Mãe d'água*, and 53: *São José*.

complex. In addition, several hybrid species have been obtained experimentally [24]. This interspecific crossing can be decisive in originating and diversifying wild species, resulting in important epidemiological consequences due to differential competence and the capacity of hybrid vectors [3, 10, 11]. Therefore, the characterization (or identification) of Triatominae specimens based only on morphological and chromatic patterns, the most common identification method, is more complex than previously believed. Studies on immature stages are crucial for group systematics. However, literature reports on immature forms of certain groups are scarce, difficult to use, or nonexistent. Many species

undergo changes in color, structure, and morphology during their development, hindering their identification [25, 26]. Triatominae nymphs at this development stage are difficult to identify. To that end, molecular analyses are successfully used to characterize morphotypes of species complexes such as *T. brasiliensis* sp., which exhibits wide chromatic and morphological variation [7, 8].

Marked differences in color pattern and ecological features among species from the *Triatoma brasiliensis* complex were detected by microsatellites, mitochondrial *12S*, and *cytochrome b* genes, reinforcing species diagnosis [7, 16]. However, individuals from subspecies *T. b. brasiliensis* and *T. b. macromelasoma* might be clustered within the same *COI* clade, since earlier studies have shown that some *cytochrome b* haplotypes of *T. b. macromelasoma* are similar to those of *T. b. brasiliensis* [16]. Moreover, these subspecies produce fertile hybrids when crossed in laboratory [3].

The basal clade of the Maximum Likelihood tree (Figure 1) was identified as subspecies *T. b. macromelasoma* because it is highly divergent (6%, as shown in the Nucleotide Distance Matrix, Table 2) from the other populations and was collected in its type locality [16]. On the other hand, all populations in the second clade were identified as *T. b. brasiliensis*. Their genetic similarity may be related to geographic proximity and similar habitat conditions. However, interpopulation divergence values (<1%) suggest that *T. b. brasiliensis* is still diversifying and/or exhibiting ongoing gene flow, probably due to human-assisted dispersal. Based on wing morphometry, Costa et al. [11] recently formulated a hypothesis that *T. b. macromelasoma* is the result of homoploid hybridization between *T. b. brasiliensis* and *T. juazeirensis* in the state of Pernambuco, and that this is a form of speciation in sympatric populations.



In northeastern Brazil, the epidemiological importance of Triatominae bugs is mainly defined by their high rate of natural *T. cruzi* infection and ability to adapt to multiple ecotopes. Control measures therefore require a precise identification of which species of the *T. brasiliensis* complex is being targeted. Moreover, it is important to understand the ecoepidemiology of Triatominae since these vectors are found in large numbers in their natural habitat [27]. In this respect, the PCR-RFLP protocol described here is suggested as rapid, relatively simple, and economical assay to distinguish *Triatoma b. macromelasoma* from *Triatoma b. brasiliensis* subspecies. Even at small geographic scales, domestic populations are genetically structured by ecological parameters, thereby exhibiting small differences from the wild counterparts from which they are derived [8]. The present study highlights the effectiveness of the *COI* gene in identifying subspecies of the *T. brasiliensis* complex and its contribution to classic taxonomy.

## Disclosure

No competing financial interests exist in the paper.

## Acknowledgments

The authors thank Dr. Maria Helena Matté (FSP/USP) for providing the restriction enzymes. Financial support was provided by *Fundação de Amparo à Pesquisa do Estado de São Paulo* (FAPESP). Daniel Pagotto Vendrami is a Master's student sponsored by FAPESP (2010/02960-3).

## References

- Á. Moncayo and A. C. Silveira, "Current epidemiological trends for Chagas disease in Latin America and future challenges in epidemiology, surveillance and health policy," *Memorias do Instituto Oswaldo Cruz*, vol. 104, no. 1, pp. 17–30, 2009.
- A. C. Silveira and M. C. Vinhaes, "Elimination of vector-borne transmission of chagas disease," *Memorias do Instituto Oswaldo Cruz*, vol. 94, no. 1, pp. 405–411, 1999.
- J. Costa, C. E. Almeida, J. P. Dujardin, and C. B. Beard, "Crossing experiments detect genetic incompatibility among populations of *Triatoma brasiliensis* Neiva, 1911 (Heteroptera, Reduviidae, Triatominae)," *Memorias do Instituto Oswaldo Cruz*, vol. 98, no. 5, pp. 637–639, 2003.
- V. J. Mendonça, M. T. A. Da Silva, R. F. De Araújo et al., "Phylogeny of *Triatoma sherlocki* (Hemiptera: Reduviidae: Triatominae) inferred from two mitochondrial genes suggests its location within the *Triatoma brasiliensis* complex," *American Journal of Tropical Medicine and Hygiene*, vol. 81, no. 5, pp. 858–864, 2009.
- H. Lent and P. Wygodzinsky, "Revision of the Triatominae (Hemiptera, Reduviidae), and their significance as vectors of Chagas' disease," *Bulletin of the American Museum of Natural History*, vol. 16, pp. 123–520, 1979.
- C. Galvão, F. M. McAloon, D. S. Rocha, C. W. Schaefer, J. Patterson, and J. Jurberg, "Description of eggs and nymphs of *Linshcosteus karupus* (Hemiptera: Reduviidae: Triatominae)," *Annals of the Entomological Society of America*, vol. 98, no. 6, pp. 861–872, 2005.
- J. Costa, M. G. R. Freitas-Sibajev, V. Marchon-Silva, M. Q. Pires, and R. S. Pacheco, "Isoenzymes Detect Variation in Populations of *Triatoma brasiliensis* (Hemiptera: Reduviidae: Triatominae)," *Memorias do Instituto Oswaldo Cruz*, vol. 92, no. 4, pp. 459–464, 1997.
- C. E. Almeida, R. S. Pacheco, K. Haag, S. Dupas, E. M. Dotson, and J. Costa, "Inferring from the Cyt B gene the *Triatoma brasiliensis* Neiva, 1911 (Hemiptera: Reduviidae: Triatominae) genetic structure and domiciliary infestation in the State of Paraíba, Brazil," *American Journal of Tropical Medicine and Hygiene*, vol. 78, no. 5, pp. 791–802, 2008.
- E. C. Borges, A. J. Romanha, and L. Diotaiuti, "Use of random amplified polymorphic DNA (RAPD) in the populational study of *Triatoma brasiliensis* Neiva, 1911," *Cadernos de Saúde Pública*, vol. 16, pp. 97–100, 2000.
- J. Costa, C. E. Almeida, E. M. Dotson et al., "The epidemiologic importance of *Triatoma brasiliensis* as a chagas disease vector in Brazil: a revision of domiciliary captures during 1993–1999," *Memorias do Instituto Oswaldo Cruz*, vol. 98, no. 4, pp. 443–449, 2003.
- J. Costa, A. T. Peterson, and J. P. Dujardin, "Morphological evidence suggests homoploid hybridization as a possible mode of speciation in the Triatominae (Hemiptera, Heteroptera, Reduviidae)," *Infection, Genetics and Evolution*, vol. 9, no. 2, pp. 263–270, 2009.
- O. Folmer, M. Black, W. Hoeh, R. Lutz, and R. Vrijenhoek, "DNA primers for amplification of mitochondrial cytochrome c oxidase subunit I from diverse metazoan invertebrates," *Molecular marine biology and biotechnology*, vol. 3, no. 5, pp. 294–299, 1994.
- J. D. Thompson, D. G. Higgins, and T. J. Gibson, "CLUSTAL W: improving the sensitivity of progressive multiple sequence alignment through sequence weighting, position-specific gap penalties and weight matrix choice," *Nucleic Acids Research*, vol. 22, no. 22, pp. 4673–4680, 1994.
- K. Tamura, D. Peterson, N. Peterson, G. Stecher, M. Nei, and S. Kumar, "MEGA5: molecular evolutionary genetics analysis using maximum likelihood, evolutionary distance, and maximum parsimony methods," *Molecular Biology and Evolution*, vol. 28, no. 10, pp. 2731–2739, 2011.
- T. Vincze, J. Posfai, and R. J. Roberts, "NEBcutter: a program to cleave DNA with restriction enzymes," *Nucleic Acids Research*, vol. 31, no. 13, pp. 3688–3691, 2003.
- F. A. Monteiro, M. J. Donnelly, C. B. Beard, and J. Costa, "Nested clade and phylogeographic analyses of the Chagas disease vector *Triatoma brasiliensis* in Northeast Brazil," *Molecular Phylogenetics and Evolution*, vol. 32, no. 1, pp. 46–56, 2004.
- C. J. Belisário, G. C. D. Pessoa, and L. Diotaiuti, "Biological aspects of crosses between *Triatoma maculata* (Erichson, 1848) and *Triatoma pseudomaculata* Corrêa & Espínola, 1964 (Hemiptera: Reduviidae)," *Memorias do Instituto Oswaldo Cruz*, vol. 102, no. 4, pp. 517–521, 2007.
- S. M. dos Santos, C. M. Lopes, J. P. Dujardin et al., "Evolutionary relationships based on genetic and phenetic characters between *Triatoma maculata*, *Triatoma pseudomaculata* and morphologically related species (Reduviidae: Triatominae)," *Infection, Genetics and Evolution*, vol. 7, no. 4, pp. 469–475, 2007.
- F. A. Carvalho-Costa, S. M. Dos Santos, M. Q. Pires, C. M. Lopes, F. Noireau, and R. S. Pacheco, "Sylvatic and peridomestic populations of *Triatoma pseudomaculata* are not significantly structured by habitat, as revealed by two genetic markers," *Journal of Vector Ecology*, vol. 35, no. 2, pp. 295–300, 2010.

- [20] A. Marcilla, M. D. Bargues, F. Abad-Franch et al., "Nuclear rDNA ITS-2 sequences reveal polyphyly of *Panstrongylus* species (Hemiptera: Reduviidae: Triatominae), vectors of *Trypanosoma cruzi*," *Infection, Genetics and Evolution*, vol. 1, no. 3, pp. 225–235, 2002.
- [21] R. P. Crossa, M. Hernández, M. N. Caraccio et al., "Chromosomal evolution trends of the genus *Panstrongylus* (Hemiptera, Reduviidae), vectors of Chagas disease," *Infection, Genetics and Evolution*, vol. 2, no. 1, pp. 47–56, 2002.
- [22] F. B. S. Dias, A. S. D. Paula, C. J. Belisário et al., "Influence of the palm tree species on the variability of *Rhodnius nasutus* Stål, 1859 (Hemiptera, Reduviidae, Triatominae)," *Infection, Genetics and Evolution*, vol. 11, no. 5, pp. 869–877, 2011.
- [23] C. E. Almeida, R. S. Pacheco, F. Noireau, and J. Costa, "*Triatoma rubrovaria* (Blanchard, 1843) (Hemiptera: Reduviidae) I: isoenzymatic and chromatic patterns of five populations from the State of Rio Grande do Sul, Brazil," *Memorias do Instituto Oswaldo Cruz*, vol. 97, no. 6, pp. 829–834, 2002.
- [24] N. Correia, C. E. Almeida, V. Lima-Neiva et al., "Cross-mating experiments detect reproductive compatibility between *Triatoma sherlocki* and other members of the *Triatoma brasiliensis* species complex," *Acta Tropica*, vol. 128, pp. 162–167, 2013.
- [25] F. T. van Emden, "The taxonomic significance of the characters of immature insects," *Annual Review of Entomology*, vol. 2, pp. 91–106, 1957.
- [26] J. Costa, A. M. Argolo, and M. Felix, "Taken of *Triatoma melanica* Neiva & Lent, 1941, new status (Hemiptera: Reduviidae: Triatominae)," *Zootaxa*, vol. 1385, pp. 47–52, 2006.
- [27] J. M. Ramsey and C. J. Schofield, "Control of Chagas disease vectors," *Salud Publica de Mexico*, vol. 45, no. 2, pp. 123–128, 2003.