Running Title: Malaria vectors in the Atlantic Forest

Ecological characterization and infection of Anophelines (Diptera: Culicidae) of the Atlantic Forest in the southeast of Brazil over a 10 year period: Has the behaviour of the autochthonous malaria vector changed?

Julyana Cerqueira Buery[1], Helder Ricas Rezende[2], Licia Natal[3], Leonardo Santana da Silva[1], Regiane Maria Tironi de Menezes[4], Blima Fux[1], Rosely dos Santos Malafronte[3], Aloisio Falqueto[1], Crispim Cerutti Junior[1].

[1]Unidade de Medicina Tropical, Universidade Federal do Espírito Santo (UFES) - Vitória (ES); [2]Secretaria de Estado da Saúde do Espírito Santo - Vitória (ES); [3]Instituto de Medicina Tropical de São Paulo, Universidade de São Paulo (USP) - São Paulo (SP); [4]Superintendencia de Controle de Endemias - São Paulo (SP).

Crispim Cerutti Junior; Av. Marechal Campos, 1 468, Maruípe, Vitória - ES, Zip Code: 29040-091, Brazil; fil.cris@terra.com.br.

ABSTRACT

In the south and southeast of Brazil, autochthonous malaria cases can be found near Atlantic Forest fragments. The transmission is not totally clarified; thus, the behaviour of the possible vectors in those regions must be observed. An entomological and natural infection study was performed on anophelines (Diptera: Culicidae) captured in the municipalities of the mountainous region of Espírito Santo state in 2004-2005. Similarly, between the years 2014 and 2015, 12 monthly collections were performed at the permanent trapping station of the study mentioned above (Valsugana Velha, Santa Teresa, ES). Light traps with CO_2 (CO_2-baited Center for Disease Control [CDC] traps) were set in open areas, at the edge of the forest (canopy and ground) and inside the forest (canopy and ground), whereas Shannon traps were set on the edge of the forest. A total of 1,414 anophelines were collected from 13 species. *Anopheles (Kerteszia) cruzii* Dyar and Knab remained the most captured species in the CO_2-baited CDC traps set in the forest canopy and was also the vector with the highest prevalence of *Plasmodium vivax* infection according to molecular PCR techniques. Regarding mosquitoes of the subgenus *Nyssorhynchus*, *P. vivax* was found only in abdomens, weakening the hypothesis that this subgenus also plays a role in malaria transmission in this specific region.

Keywords: Malaria, *Plasmodium vixax*, Anopheles.

Sponsorship: Espírito Santo Research Foundation (Fundação de Amparo à Pesquisa e Inovação do Espírito Santo – FAPES).

INTRODUCTION

Despite being a highly prevalent disease in the Amazon region, malaria remains residual in Atlantic Forests of the south and southeast regions of Brazil. In this region, the disease is known as bromeliad-malaria since the vectors of the genus *Anopheles* reproduce in the whorls of *Bromeliaceae,* which are typical plants of this biome (Downs & Pittendrigh, 1946). The presence of a few autochthonous (sometimes asymptomatic) human cases, the spatial distance between the cases and the low parasitemia by microscopy weaken the possibility of the traditional transmission chain. It is believed that the cycle of *Plasmodium spp.* is maintained by reservoirs of the parasite, in the forest or in the rural population, represented by both apes and humans, asymptomatic or not. Studies have been conducted to clarify these beliefs (Curado et al., 1997; Duarte et al., 2006; Cerutti et al., 2007; Rezende et al., 2009; Meneguzzi et al., 2009). Regarding the vector, in the state of Espírito Santo, Brazil, 26 species of *Anopheles sp.* have been identified, indicating an abundance of anophelines (Coutinho, 1947; Andrade&Brandão, 1957; Ferreira et al., 1964; Natal, 2007; Sallum, 2008; Rezende et al. 2009; Meneguzzi et al. 2009; Silva et al., 2013). The vectors of greater vectorial capacity and competence belong to the subgenus *Kerteszia*, mainly *Anopheles cruzii*, in the south and southeast Brazilian states (Meneguzzi et al., 2009). However, species of anophelines involved in the dynamics of malaria transmission outside the Amazon region vary according to environmental and epidemiological conditions (Pina-Costa et al., 2014). The autochthonous cases of extra-Amazonian malaria are found in mild mountainous regions and are associated with agricultural activities near the forest that are performed by young men (Cerutti Jr et al., 2007). States such as São Paulo, Santa Catarina and Espírito Santo are covered by dense Atlantic Forest regions (IESB, 2007). Among them, Espírito Santo has recorded the largest number of bromeliad-malaria cases in recent

years. The biome in question is very humid, with abundant rainfall and vegetation that favours the reproduction of the anophelines. In addition, bromeliad-malaria is beginning to be used as a biological marker of human activities within these forests (Gomes, 1985; Rezende et al., 2013). Rezende et al. (2009) found anopheline behaviour similar to that expected, based on the literature, in Valsugana Velha, Santa Teresa municipality: a high prevalence of *A. cruzii* was reported in a mountainous region of the Atlantic Forest, in an endemic area of the disease in the state of Espírito Santo. However, considering the hypothesis of a typical transmission model of zoonosis in this scenario, this model would require a stable and constant vectorial behaviour. Such behaviour, therefore, must be re-evaluated to establish transmission characteristics over the years.

MATERIALS AND METHODS

Description of the study site. The study was performed in the rural area of the municipality of Santa Teresa, which is located approximately 78 km from Vitória, in the state of Espírito Santo (ES), Brazil. The permanent trapping station is in Valsugana Velha (19°57'58.4 "S, 40°34'45.2" W), where cases of residual malaria from Atlantic Forest systems were recorded in the study by Cerutti Jr. et al. (2007) (Figure 1).

[Figure 1. Political map of the municipality of Santa Teresa, highlighting the rural community where the collections were performed. Source: Instituto Jones dos Santos Neves, 2013.]

Collection of anophelines. Hourly collections of anophelines were conducted at a permanent trapping station located in the bromeliad-malaria transmission area. These

collections were performed one day each month for one year, from June 2014 to May 2015, totalling 12 collections. Two capture methods were used:

- Center for Disease Control (CDC) light traps with CO_2 (Gomes, 1985) that were set in open areas (peridomiciliary environment = PD) , at the edge of the forest (canopy and ground) and inside the forest (canopy and ground) and
- Shannon traps (Bustamante, 1951), set at the edge of the forest.

The five CDC traps were set simultaneously, two of them at 15 metres from the ground in the canopy (on the edge and inside the forest), two at one metre from the ground (at the edge and inside the forest), and one at the border between the forest and the area close to the dwellings. The collections lasted for 12 hours, with the traps placed at night (6:00 PM) and removed in the morning (6:00 AM). For the Shannon traps, the insects were captured during the first 4 hours after sunset (6:00 PM to 10:00 PM) each month.

Storage and identification of insects. The specimens were stored in tubes containing isopropanol and later identified using the identification keys proposed by Consoli&Lourenço-de-Oliveira (1994). The identification was made by a team from the Entomology and Malacology Centre of Espírito Santo (Núcleo de Entomologia e Malacologia do Espírito Santo - NEMES/ES).

Molecular techniques. The DNA for the detection of *Plasmodium sp.* was obtained from the thorax, abdomen or entire mosquito of the mosquito groupings in pools (maximum of 10 samples/pool), depending on the subgenus of the specimens. Those of the subgenus *Nyssorhynchus* were sectioned, whereas those of the subgenus *Kerteszia* were processed in whole. The same extraction kit (DNeasy Blood & Tissue Kit, Qiagen, Germany) was used, following the manufacturer's instructions. Each pool included females of the same species, collected in the same type of trap, on the same date. The

presence of *P. vivax* or *P. malariae* in the pools was detected using the nested-PCR protocol described by Kimura et al. (1997) and modified by Win et al. (2002). The target was the gene encoding the 18S ribosomal subunit. The amplification products were analysed by electrophoresis in 2% agarose gel under ultraviolet light. In positive cases, 100-bp fragments were amplified.

Statistical analysis. To determine the importance and distribution of the various *Anopheles* species, analyses of diversity, dominance and abundance were performed using Shannon's diversity index (H') and Simpson's dominance index (D). In the various comparisons, the level of significance was set at 5%. Bivariate Spearman correlation calculations were used to determine the relationship between anopheline capture, temperature and rainfall (data provided by the Capixaba Institute of Research, Technical Assistance and Rural Extension [Instituto Capixaba de Pesquisa, Assistência Técnica e Extensão Rural - INCAPER]), also at a level of 5%.

ETHICAL STANDARDS

During the collection process, no harm was inflicted to the environment. The team members wore long clothing, gloves and hats with head nets to avoid anopheline mosquito bites. Boots were also used to prevent bites by venomous animals. A license to capture arthropod insects was obtained via the SISBIO/Chico Mendes Institute (ICMBio/IBAMA/MMA) under number 19227-1.

RESULTS

A total of 1,414 specimens were captured, resulting in a set of 13 species. The largest number of specimens was collected in April 2015 (341) and the smallest number of

specimens in March 2015 (05). In all collections, there was a predominance of *Anopheles (K.) cruzii*, totalling 1,044 of the 1,414 mosquitoes captured (Table 1). The trap in which most mosquitoes were captured was the CO_2-baited CDC trap placed in the canopy, in which *A. cruzii* was also predominant (Graph 1).

[Graph 1. Percentage of species captured per trap, from June 2014 to May 2015, at the permanent trapping station in Santa Teresa, ES.]

Climate. The months of November 2014, September 2014 and April 2015 showed the highest capture frequencies (Graph 2). There was no simultaneous capture of all species in the monthly collections, and all species were absent for at least 1 month during the total collection period. *A. cruzii* was not captured only in June 2014 and March 2015 (Graph 2). In fact, those were the only months in which *A. cruzii* ceased to be the predominant species and was replaced with *A. strodei* and *A. triannulatus*, respectively.

[Graph 2. Species captured each collection month, from June 2014 to May 2015, at the permanent trapping station of Santa Teresa, ES.]

A negative trend was observed for the correlation between the number of anophelines of the subgenus *Nyssorhynchus* and the temperature, although without statistical significance ($r = -0.04$, $p = 0.89$). Additionally, a positive trend was observed for the correlation between the number of anophelines and rainfall ($r = 0.17$, $p = 0.59$) but also with no statistical significance. For the subgenus *Kerteszia*, the correlation between capture frequency and temperature ($r = -0.004$, $p = 0.99$) or rainfall ($r = -0.13$, $p = 0.68$)

revealed negative coefficients but with no statistical significance. The monthly average temperature and the rainfall in Valsugana Velha are shown in Graph 3.

[Graph 3. Temperature and rainfall data in Santa Teresa, ES, between June 2014 and May 2015.]

Spatial distribution. *A. strodei* and *A. triannulatus* were more frequently captured in the Shannon traps, and *A. cruzii* was captured more frequently in the CO_2-baited CDC traps (Graph 1) located in the tree canopy. Simpson's dominance index (D) reveals that dominance in the Shannon trap (D1 = 0.227) was greater than the dominance of individuals collected in the CO_2-baited CDC trap (D2 = 0.172) ($p < 0.02$), both at the forest edge.

Based on Shannon's diversity index (H'), the diversity of anophelines collected in the Shannon trap (H'2 = 1.866) was higher compared to that of the anophelines collected in the CDC trap at the forest edge, near the dwellings (H'1 = 1.734) ($p = 0.004$). *Anopheles strodei* was captured in larger numbers at forest edge areas, followed by *A. triannulatus*. *A. lanei* and *A. rangeli* were captured only in the CO_2-baited CDC trap located near the dwellings, whereas *A. homunculus* was only captured in the CDC traps set in the tree canopy.

Natural infection rates (Table 2). Of the total pool of specimens investigated, 13 were positive for *Plasmodium vivax* (Figure 2). Of these, 10 belonged to the species *A. cruzii*, and three were abdomens that belonged to specimens of the subgenus *Nyssorhynchus*. As shown in Table 2, most of the infected mosquitoes were *A. (K.) cruzii*, captured in CO_2-baited CDC traps located in the tree canopy. Graph 4 shows the monthly

distribution of captured mosquitoes in which *P. vivax* was found. *P. malariae* infection was not detected.

[Figure 2. Ethidium bromide-stained agarose gel, photographed using the Alpha Imager software, with 13 samples of pools of *P. vivax*-positive anopheline females. Source: Buery JC and Natal L, 2016.]

[Graph 4. Chronological distribution of *P. vivax*-infected anopheline females captured in the permanent trapping station in Santa Teresa, ES.]

DISCUSSION

Bromeliad malaria became more evident in the Brazilian south and southeast Atlantic forest regions after control measures implemented by the government eliminated transmission in flat areas exploited by humans (Ministério da Saúde (BR), 2015; Barata, 1998). As residual foci remained in the eradication areas, bromeliad-malaria or malaria of Atlantic Forest systems began to be studied with more intensity in the 1980s and 1990s (Duarte et al., 2013). With a more consistent understanding of vector ecology and outbreak epidemiology, more evidence has emerged in favour of behaviour consistent with a zoonosis. The zoonosis hypothesis presupposes the concomitant existence of simian malaria. Such an existence has been proven in the past (Duarte et al., 2008), but its prospective evaluation demands extremely complex logistics. Therefore, the evaluation of vector behaviour favourable to enzootic transmission is more feasible and may provide indirect evidence of the presence of the infection among monkeys and of its source of occurrence in human cases. Given the need for stable and constant vector behaviour for the maintenance of simian malaria, follow-up studies of the endemic

221 regions are necessary. In addition, according to Marrelli et al. (2007), the entire Atlantic
222 Forest territory that survived deforestation must be carefully monitored due to
223 environmental changes and the possibility of maintaining the infection cycle given the
224 high number of asymptomatic individuals who can act as reservoirs of the parasite in
225 those regions.

226 In the present study, *A. cruzii* still prevailed as the main vector found at the Valsugana
227 Velha trapping station. Between 2004 and 2005 (Rezende et al. 2009), 61.2% of the
228 2,290 mosquitoes collected belonged to this species. In 2014 and 2015, 73.8% of the
229 1,414 anophelines captured were *A. cruzii*. Therefore, an increase in the proportion of
230 the main vector of bromeliad malaria was observed in the local fauna. Older studies
231 such as those of Deane (1988) and more recent studies such as those of Duarte et al.
232 (2013), Neves et al. (2013) and Kirchgatter et al. (2014) have previously demonstrated
233 the magnitude of the presence of this species in Atlantic Forest regions with preserved
234 native forest.

235 The distribution of *A. cruzii* when moving from the inside to the edge of the forest has
236 also remained the same over time. As Rezende et al. (2009) described in 2004 and 2005,
237 the anophelines of this species appeared mostly in the tree canopy inside the forest,
238 whereas the numbers decreased drastically in the traps that were set closer to the forest
239 edge and the dwellings. The anopheline fauna was dominated by species such as *A.*
240 *strodei* and *A. triannulatus* closer to human-occupied areas. In São Paulo, between 2009
241 and 2011, this trend was also detected when comparing anthropic and wild areas in the
242 town of Parelheiros. There, 438 *A. cruzii* were recorded in a certain anthropic area and
243 4,832 in the wild area (Duarte et al., 2013).

The absolute predominance of *A. cruzii* in the canopy, observed in the present study, indicates a non-synanthropic behaviour (Forattini et al., 1990) and reinforces its role in the transmission of simian malaria. However, despite being present mostly in the canopy, some specimens were found in the CO_2-baited CDC traps set in the ground and in the Shannon traps, which points to the possibility of these vectors descending from their preferential site in the canopy, at which time they could incidentally transmit the parasites to humans. The presence of these specimens reinforces the hypothesis that bromeliad-malaria is maintained in the region by means of simian infection. The fact that humans are in the forest and that *A. cruzii* comes down from the canopy to feed creates the conditions for *Plasmodium spp.* originating from the monkeys in the canopy to infect humans.

The morphometric diagnosis studies of Lorenz et al. (2012) showed a differentiation between *A. cruzii* and *A. homunculus* not predicted in the study of Rezende et al. (2009). However, only 39 *A. homunculus* were captured during the collection period between 2014 and 2015, against 1,045 duly identified *A. cruzii*.

Unexpectedly, the season during which most mosquitoes were captured was not the summer. Interestingly, the seasons with milder weather had the largest capture rates. For example, the most successful captures occurred in the months of September 2014 and April 2015. This result corroborates the study of Rezende et al. (2009), who suggested the adaptation of anophelines to mild environments. There was also a high number of specimens collected in November 2014. The mean rainfall at that time was the highest of that year (mean of 207.6 mm/month), and the temperature was mild on collection day (ranging from 19.7 to 21.3 °C). That month, the "rain" factor may have triggered an increase in the mosquito population.

In this study, 13 pools of mosquitoes were positive for *P. vivax*. In 2004 and 2005, 10 pools of anophelines infected by the same species of the parasite were obtained (Rezende et al., 2009). However, unlike 10 years ago, the PCR reactions for the 2014 and 2015 collections did not show infection in the thorax of mosquitoes of the subgenus *Nyssorhynchus*. The infective form should reach the salivary gland of the anopheline for infection to occur. Thus, the separation into thorax and abdomen during the experiments reinforces evidence of the participation of other vector species in the transmission chain. Because *Kerteszia* was a subgenus of a known vector, there was no separation of the body to perform the experiments, as occurred in mosquitoes of the subgenus *Nyssorhynchus*, to assess the possibility that these mosquitoes participate in the transmission chain. Since no infection was detected in the *Nyssorhynchus* thoraxes, unlike 10 years ago, *Nyssorhynchus* may have stopped playing the role of secondary vector and *A. cruzii* may currently be the only bromeliad-malaria vector in this area. The progressive exploitation of the rural and forest environment by the local inhabitants may have led to greater spatial distances between the transmission events and the anthropic environment, where *Nyssorhynchus* has greater dominance. In these conditions, given the *A. cruzii* dominance, *Nyssorhynchus* does not have the opportunity to become infected. Humans venture into the forest to clean the river springs or to gather firewood and thus acquire the disease. Once they return to their homes, they likely infect the *Nyssorhynchus* near their dwellings.

Infected *A. cruzii* were collected in CO_2-baited CDC traps in the tree canopy inside the forest and in one CO_2-baited CDC trap located near the ground, at a height of one metre. This fact reinforces the idea that infections can occur in both habitats as a result of the acrodendrophilic behaviour with eventual descent to lower heights, where they can feed

on non-usual hosts. This fact also reinforces the possibility of the disease being a zoonosis in regions such as Valsugana Velha, in Santa Teresa, ES.

Regarding the Shannon's dominance index (H'), there was greater dominance in the Shannon trap than in the CO_2-baited CDC trap, both at the forest edge.

Regarding Simpson's diversity index (D), the diversity of anophelines collected with the Shannon trap was also higher than that observed in the CO_2-baited CDC trap in the same habitat. The placement of the trap near the body of water, where there were *Nyssorhynchus* breeding sites, would justify the greater diversity.

These data corroborate a study conducted in the same region and published in 2013 (Rezende et al., 2013), when greater dominance and diversity of anophelines were observed in an anthropic environment with the presence of malaria. These findings reinforce the role of human occupation in the determination of both anopheline distribution and behaviour since both indices show higher values in collections performed in an environment closer to human dwellings.

The study shows that there was little change in vector behaviour in the region studied. *A. cruzii* remains the most infected anopheline in Valsugana Velha, and mosquitoes of the subgenus *Nyssorhynchus* do not appear to participate in the transmission chain. The acrodendrophilic behaviour of *A. cruzii*, particularly those infected, reinforces the hypothesis that the presence of *P. vivax* in these specimens arises from blood feeding in animals that live in the tree canopy, such as simians.

ACKNOWLEDGEMENTS

We would like to thank the State Health Department of Espírito Santo and the Entomology and Malacology Centre of Espírito Santo/SESA for the logistical support

and supply of equipment in the field; Claudiney Biral dos Santos for technical support; Filomena E C Aguiar, Creuza Rachel Vicente, Marcelo Urbano Ferreira, Priscila Thihara Rodrigues and Lais Camoese Salla for the critical reading of the manuscript; the owners of the Recanto da Preguiça farm for allowing sampling at that site; and the Capixaba Institute of Research, Technical Assistance, and Rural Extension (INCAPER/ES) for providing the rainfall and temperature data.

FINANCIAL SUPPORT

This work was supported by the Espírito Santo Research Foundation (Fundação de Amparo à Pesquisa e Inovação do Espírito Santo - FAPES) and by the Research Program for Unified Health System of Brazilian Ministry of Health (Programa de Pesquisa para o Sistema Único de Saúde do Ministério da Saúde) (Grant number 65834119/2014).

CONFLICTS OF INTEREST

None.

REFERENCES

Andrade RM, Brandão H. Contribuição para o conhecimento da fauna de anofelinos do estado do Espírito Santo: área de distribuição e incidência das espécies por cidades, vilas e povoados. Rev Bras Malariol Doenças Trop. 1957; 9: 391-403.

Barata RB. Malária e seu controle. São Paulo: Editora Hucitec; 1998.

341 Bustamante FM. Shannon dawn trap: its use in the verification of the durability of
342 residual toxic effects of insecticides. Folha Med . 1951; 32(8):53-55.
343
344 Cerutti Jr C, Boulos M, Coutinho AF, Hatab M do CLD, Falqueto A, Rezende HR,
345 et al. Epidemiologic aspects of the malaria transmission cycle in an area of very low
346 incidence in Brazil. Malar J. 2007; 6:33.
347
348 Consoli RAGB, Oliveira RL. Principais mosquitos de importância sanitária no
349 Brasil. Cad Saúde Pública. 1994; 11: 228 p.
350
351 Coutinho JO. Contribuição para o estudo da distribuição geográfica dos anofelinos
352 do Brasil (dissertação). São Paulo, SP, Brasil: Faculdade de Medicina da
353 Universidade de São Paulo; 1947.
354
355 Curado I, Duarte AMRC, Lal AA, Oliveira SG, Kloetzel JK. Antibodies anti
356 Bloodstream and Circumsporozoite Antigens (*Plasmodium vivax* and *Plasmodium*
357 *malariae*/*P. brasilianum*) in Areas of Very Low Malaria Endemicity in Brazil. Mem
358 Inst Oswaldo Cruz. 1997; 92(2):235–43.
359
360 Deane LM. Malaria studies and control in Brazil. Am J Trop Med Hyg. 1988;
361 38(2):223-230.
362 Downs WG, Pittendrigh CS. Bromelian malaria in Trinidad, British West Indies;
363 Am J Trop Med. 1946; 26: 47-66.
364

Duarte AMRC, Porto MAL, Curado I, Malafronte RS, Hoffman EHE, Oliveira SG *et al.* Widespread occurence of antibodies against Circumsporozoite protein and against blood forms of *Plasmodium vivax, P. falciparum* and *P. malariae* in Brazilian wild monkeys. Int J Primatol. 2006; 35: 87-96.

Duarte AM, Malafronte RS, Cerutti C Jr, Curado I, Paiva BR, Maeda AY et al. Natural Plasmodium infections in Brazilian wild monkeys: reservoirs for human infections? Acta Trop. 2008; 07:179–85.

Duarte AM, Pereira DM, de Paula MB, Fernandes A, Urbinatti PR, Ribeiro AF, et al. Natural infection in anopheline species and its implications for autochthonous malaria in the Atlantic Forest in Brazil. Parasit Vectors. 2013; 6:58

Ferreira E. Distribuição geográfica dos anofelinos no Brasil e sua relação com o estado atual da erradicação da malária. Rev Bras Malariol Doenças Trop. 1964; 16: 329-48.

Forattini OP, Kakitani I, Massad E, Marucc D. Studies on mosquitoes (Diptera: Culicidae) and anthropic environment. 11 - Biting activity and blood-seeking parity of Anopheles (Kerteszia) in South-Eastern Brazil. Rev Saude Publica. 1990; 24(2): 107-14.

Gomes AC, Rabello EX, Natal D. A new collecting chamber for a CDC-miniature trap. Rev Saude Publica. 1985; 19(2):190-91.

Instituto de Estudos Socioambientais do Sul da Bahia (IESB). Projeto de conservação e utilização sustentável da diversidade biológica brasileira – PROBIO. Levantamento da Cobertura Vegetal Nativa do Bioma Mata Atlântica. Rio de Janeiro: Instituto de Geociências da Universidade Federal do Rio de Janeiro, 2007.

Kimura M, Kaneco O, Liuc Q, Zhouc M, Kawamotoc F, Watayad Y, et al. Identification of the four species of human malaria parasites by nested PCR that targets variant sequences in the small subunit rRNA gene. Parasitol Int. 1997; 46:91-95.

Kirchgatter K, Tubaki RM, Malafronte RM, Alves IC, Lima GFMC, Guimarães LO et al. *Anopheles (Kerteszia) cruzii* (Diptera: Culicidae) in peridomiciliary area during asymptomatic malaria transmission in the Atlantic Forest: molecular identification of blood-meal sources indicates humans as primary intermediate hosts. Rev Inst Med Trop São Paulo. 2014; 56(5):403-09.

Lorenz C, Marques T, Sallum MA, Suesdek L. Morphometrical diagnosis of the malaria vectors *Anopheles cruzii*, *An. homunculus* and *An. bellator*. Parasit Vectors. 2012; 5(1):257.

Marrelli MT, Malafronte RS, Sallum MAM, Natal D. Kerteszia subgenus of Anopheles associated with the Brazilian Atlantic rainforest: current knowledge and future challenges. Malar J .2007; 6:127.

Meneguzzi VC, Santos CB, Pinto IS, Feitoza LR, Feitoza HN, Falqueto A. Use of geoprocessing to determine malaria risk areas and evaluation of the vectorial importance of anopheline mosquitoes (Diptera: Culicidae) in Espírito Santo, Brazil. Mem Inst Oswaldo Cruz. 2009; 104: 570-75.

Ministério da Saúde (BR). Ministério da Saúde lança Plano de Eliminação da malária no Brasil. Brasília: Ministério da Saúde, 2015.

Natal D. First record of *Anopheles* (*Anopheles*) *costai* Fonseca & Ramos, 1939 in Espírito Santo State. Rev Inst Med Trop São Paulo. 2007; 49(5):323-26.

Neves A, Urbinatti PR, Malafronte RS, Fernandes A, Paganini WS, Natal D. Malaria outside the Amazon region: Natural Plasmodium infection in anophelines collected near an indigenous village in the Vale do Rio Branco, Itanhaém, SP, Brazil. Acta Trop. 2013; 125: 102-06.

Pina-Costa A, Brasil P, Santi S MD, de Araujo MP, Suárez-Mutis MC, Santelli AC, et al. Malaria in Brazil: what happens outside the Amazonian endemic region. Mem Inst Oswaldo Cruz. 2014; 109(5):618-33.

Rezende HR, Soares RM, Cerutti C, Alves IC, Natal D, Urbinatti PR, et al. Entomological characterization and natural infection of anophelines in an area of the Atlantic Forest with autochthonous malaria cases in mountainous region of Espírito Santo State, Brazil. Neotrop Entomol. 2009; 38(2):272–80.

Rezende HR, Falqueto A, Urbinatti PR, De Menezes RM, Natal D, Cerutti C Jr. Comparative study of distribution of anopheline vectors (Diptera: Culicidae) in areas with and without malaria transmission in the highlands of an extra-Amazonian region in Brazil. J Med Entomol. 2013; 50(3):598-602.

Sallum MAM. Primeiro registro de *Anopheles* (*Kerteszia*) *homunculus* Komp (Diptera, Culicidae) no Estado do Espírito Santo, Brasil. Rev Bras Entomol. 2008; 52: 671-73.

Silva KS, Pinto IS, Leite GR, Virgens TM, Santos CB, Falqueto A. Ecology of Anopheline Mosquitoes (Diptera: Culicidae) in the Central Atlantic Forest Biodiversity Corridor, Southeastern Brazil. J Med Entomol. 2013; 50(1):24-30.

Win TT, Lin K, Mizuno S, Zhou M, Liu Q, Ferreira MU, et al. Wide distribution of *Plasmodium ovale* in Myanmar. Trop Med Int Health. 2002; 7(3):231–39.

454 **TABLES**

455 **Table 1.** Percentage of anopheline species found between June 2014 and May 2015 at
456 the permanent trapping station in Santa Teresa, ES.

Species	**Number**	**[%]**
A. (K) cruzii	1044	73.8
A. (N) strodei	103	7.3
A. (N) triannulatus	84	6.0
A. (N) evansae	52	3.7
A. (K) homunculus	39	2.7
A. (N) galvaoi	29	2.0
A. (N) lutzii	21	1.5
A. (N) albitarsis	17	1.2
A. (N) argyritarsis	09	0.6
A. (N) parvus	07	0.5
A. (A). mediopunctatus	05	0.4
A. (N) rangeli	03	0.2
A. (N) lanei	01	0.1
TOTAL	**1414**	**100**

457 **Table 2.** Species, traps and collection dates for *P. vivax*-positive mosquitoes at the

458 permanent trapping season in Santa Teresa, ES.

Sample	Species	Trap	Date
260a*	*A. (N.) lutzii*	Shannon	September 2014
279	*A. (K.) cruzii*	CDC canopy	August 2014
343	*A. (K.) cruzii*	CDC canopy	August 2014
479	*A. (K.) cruzii*	CDC canopy	October 2014
485	*A. (K.) cruzii*	CDC canopy	October 2014
494a*	*A. (N.) evansae*	Shannon	October 2014
632	*A. (K.) cruzii*	CDC ground	November 2014
633	*A. (K.) cruzii*	CDC canopy	November 2014
868	*A. (K.) cruzii*	CDC canopy	February 2015
1142	*A. (K.) cruzii*	CDC canopy	April 2015
1144	*A. (K.) cruzii*	CDC canopy	April 2015
1248	*A. (K.) cruzii*	CDC canopy	April 2015
1294a*	*A. (N.) strodei*	CDC near dwellings	May 2015

459 *a = abdomen.

460

FIGURES

Figure 1. Political map of the municipality of Santa Teresa, highlighting the rural community where the collections were performed. Source: Instituto Jones dos Santos Neves, 2013.

Figure 2. Ethidium bromide-stained agarose gel, photographed using the Alpha Imager software, with 13 samples of pools of *P. vivax*-positive anopheline females. Source: Buery JC and Natal L, 2016.

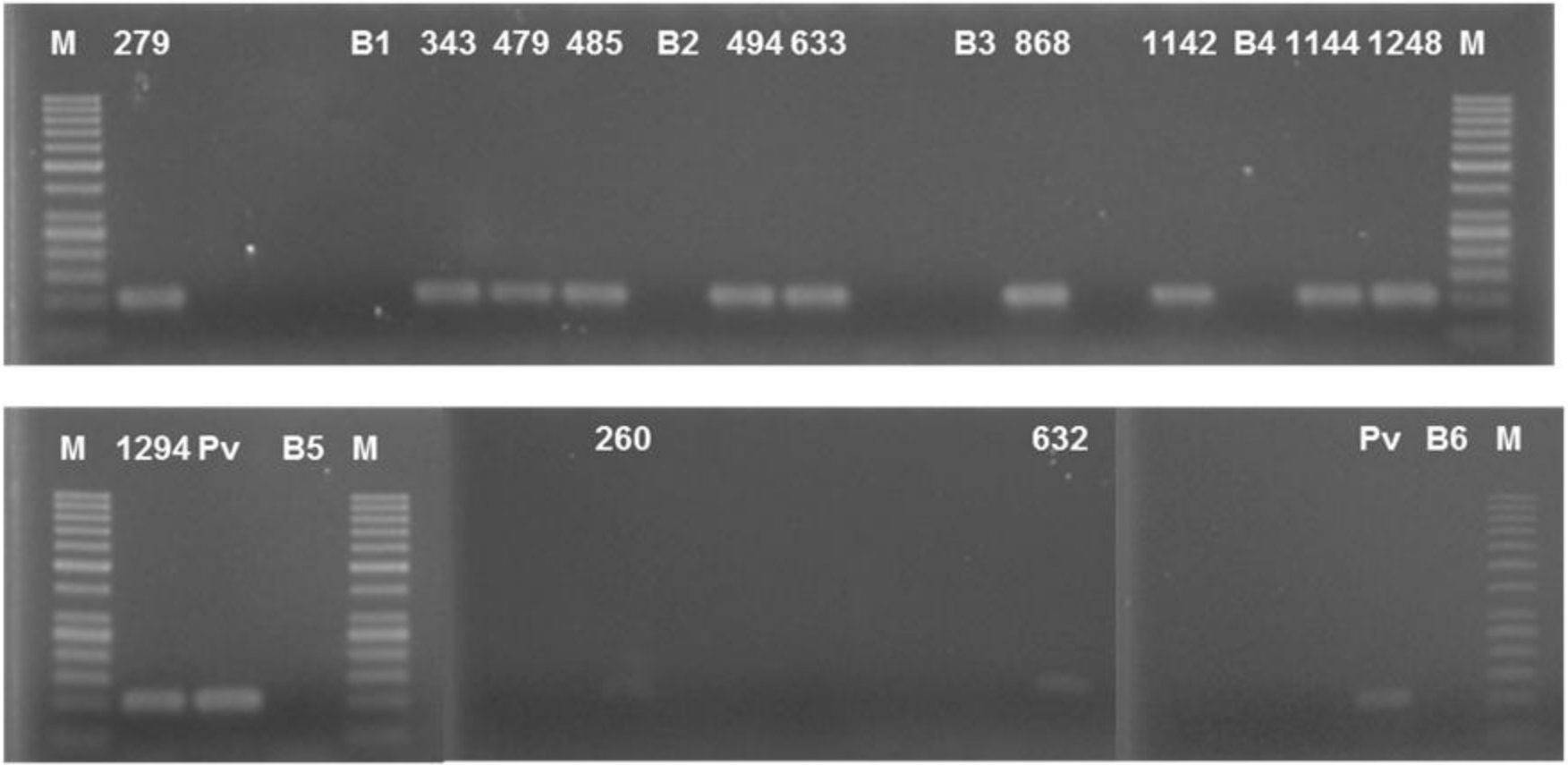
Graph 1. Percentage of species captured per trap, from June 2014 to May 2015, at the permanent trapping station in Santa Teresa, ES.

Graph 2. Species captured each collection month, from June 2014 to May 2015, at the permanent trapping station of Santa Teresa, ES.

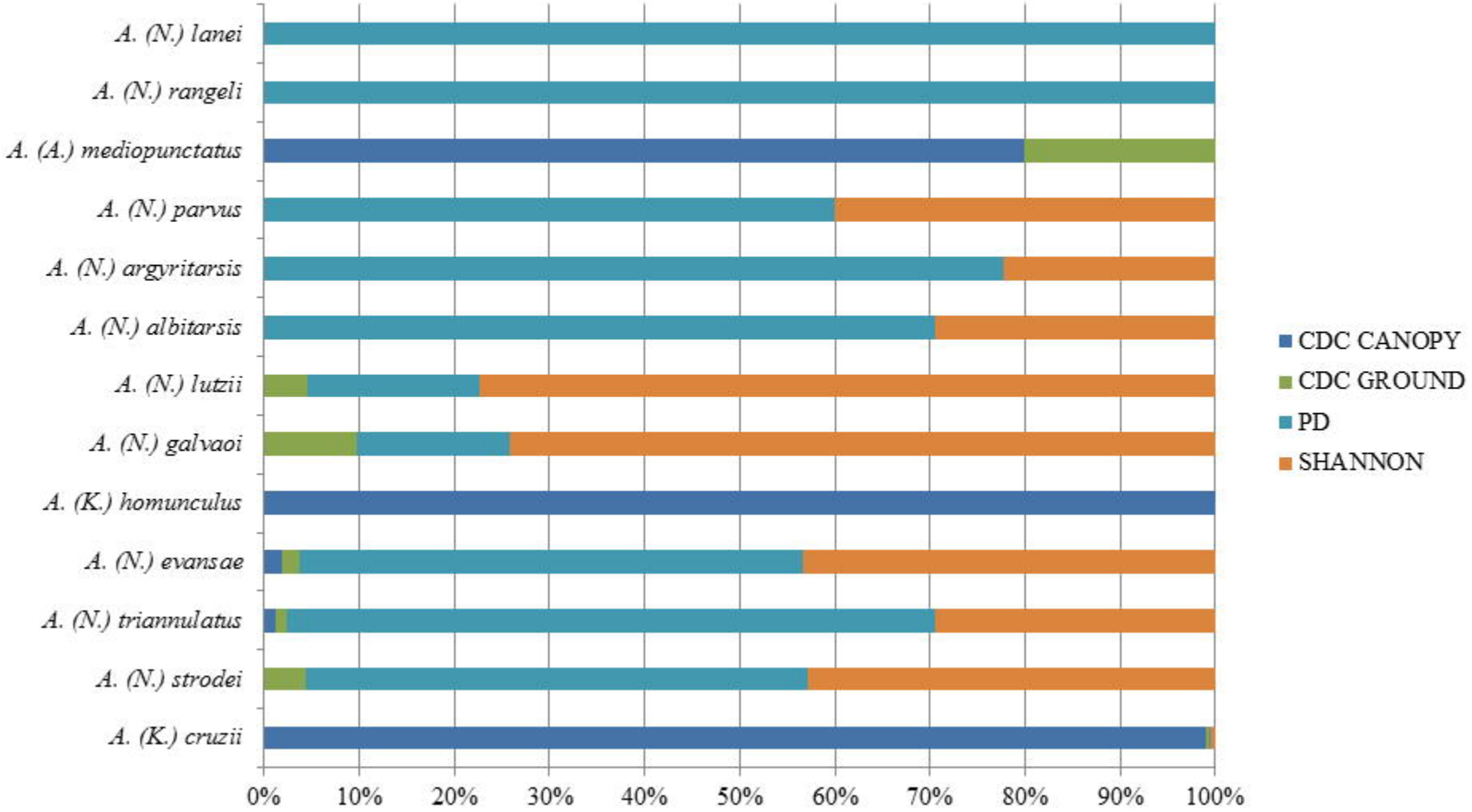
Graph 3. Temperature and rainfall data in Santa Teresa, ES, between June 2014 and May 2015.

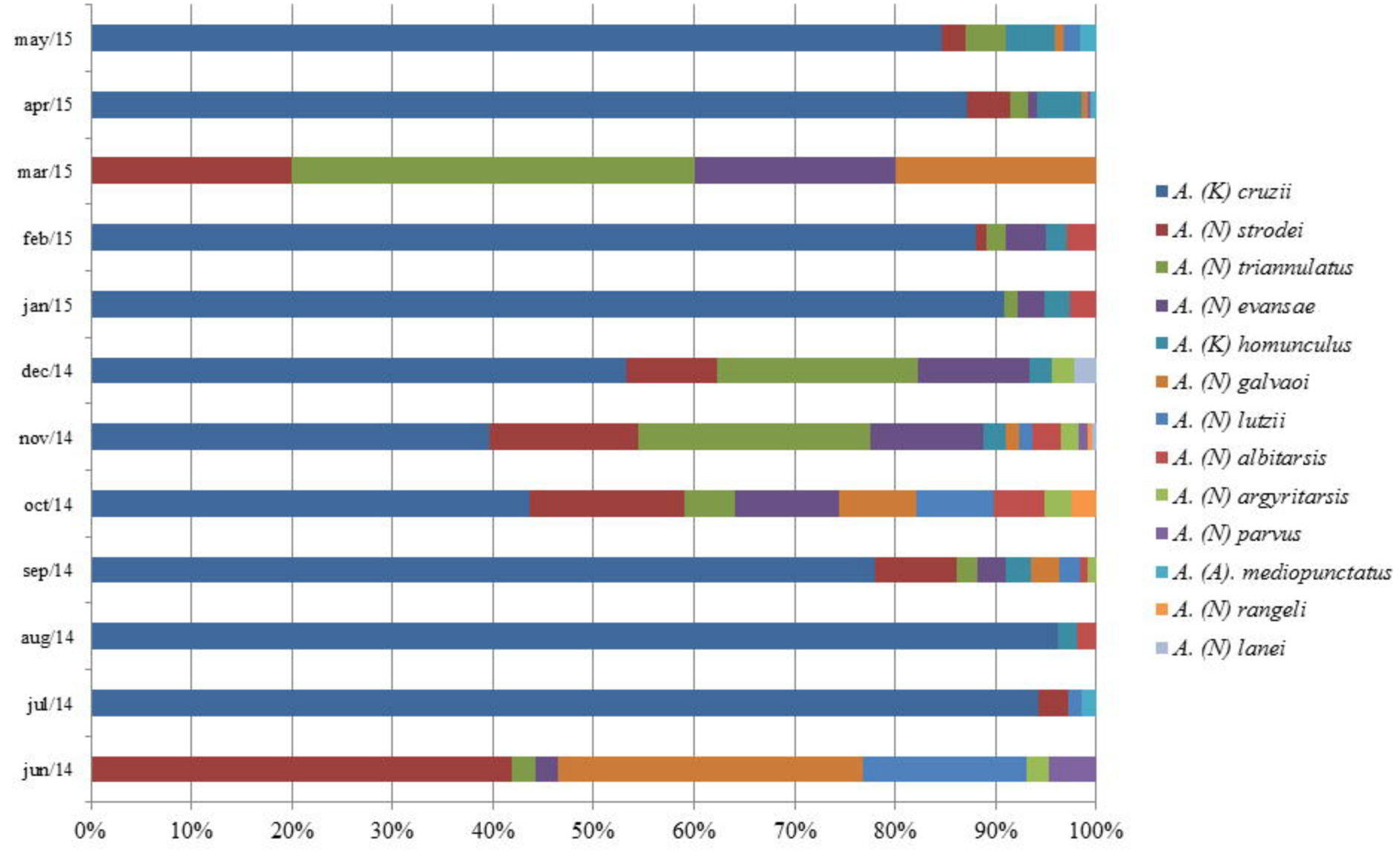
Graph 4. Chronological distribution of *P. vivax*-infected anopheline females captured in the permanent trapping station in Santa Teresa, ES.

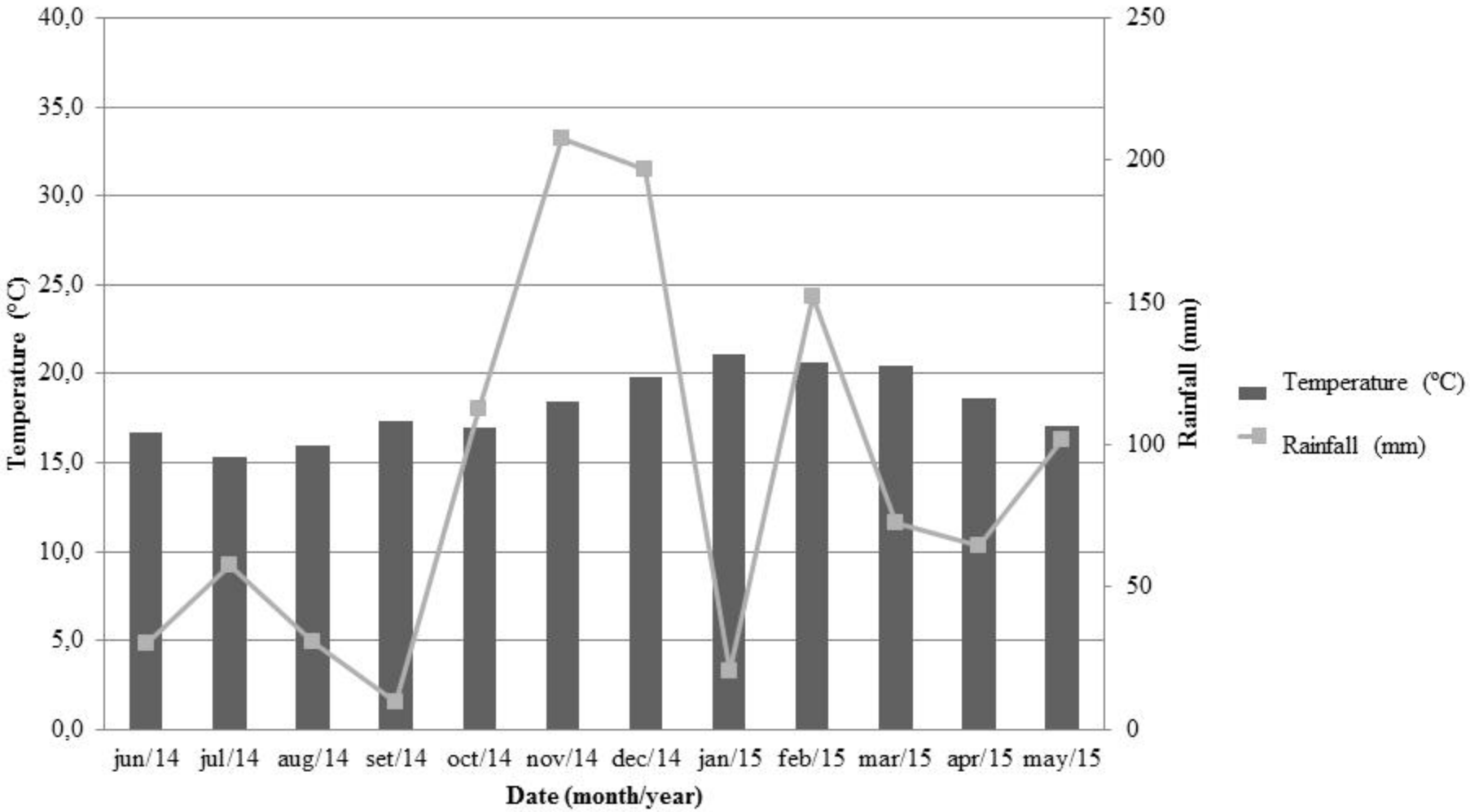

Brasil
ES
BA
MG
RJ
São Roque do Canaã
João Neiva
Rib. Alegre
Julião
Vinte e Cinco de Julho
Quinze de Agosto
Lampê
São José do Petrópolis
Córrego Seco de São João
Córrego do Mangangá
Alto Vinte e Cinco de Julho
Santo Antônio
Cab. Vinte e Cin. de Julho
Lombardia
Itaguaçu
Córrego Frio
Alto Santa Júlia
Barra do Perdido
Salto Anselmo
Ibiraçu
Itanhagá
São Paulo do Rio Perdido
Barra da Taboca
Pedra Alegre
São Sebastião de Várzea Alegre
Santo Hilário
Córrego São Martim
Alto Santo Antônio
Alto Várzea Alegre
Nova Valsugana
Rio Cinco de Novembro
Alto Goiapaba Açu
Várzea Alegre
São José do Caldeirão
Res. Biológica de Nova Lombardia
Tabocas
Goiapaba Açu
Toma Vento
Vargem Alta
Caldeirão
Rio Saltinho
Santa Teresa
Fundão
Itarana
Santa Lúcia
Valão de São Pedro
Serra dos Pregos
Cabeceira do Rio Bonito
Valsugana Velha
Alto Caldeirão
Res. Bio Augusto Ruschi
Serra do Gelo
Santa Maria de Jetibá
Santa Leopoldina

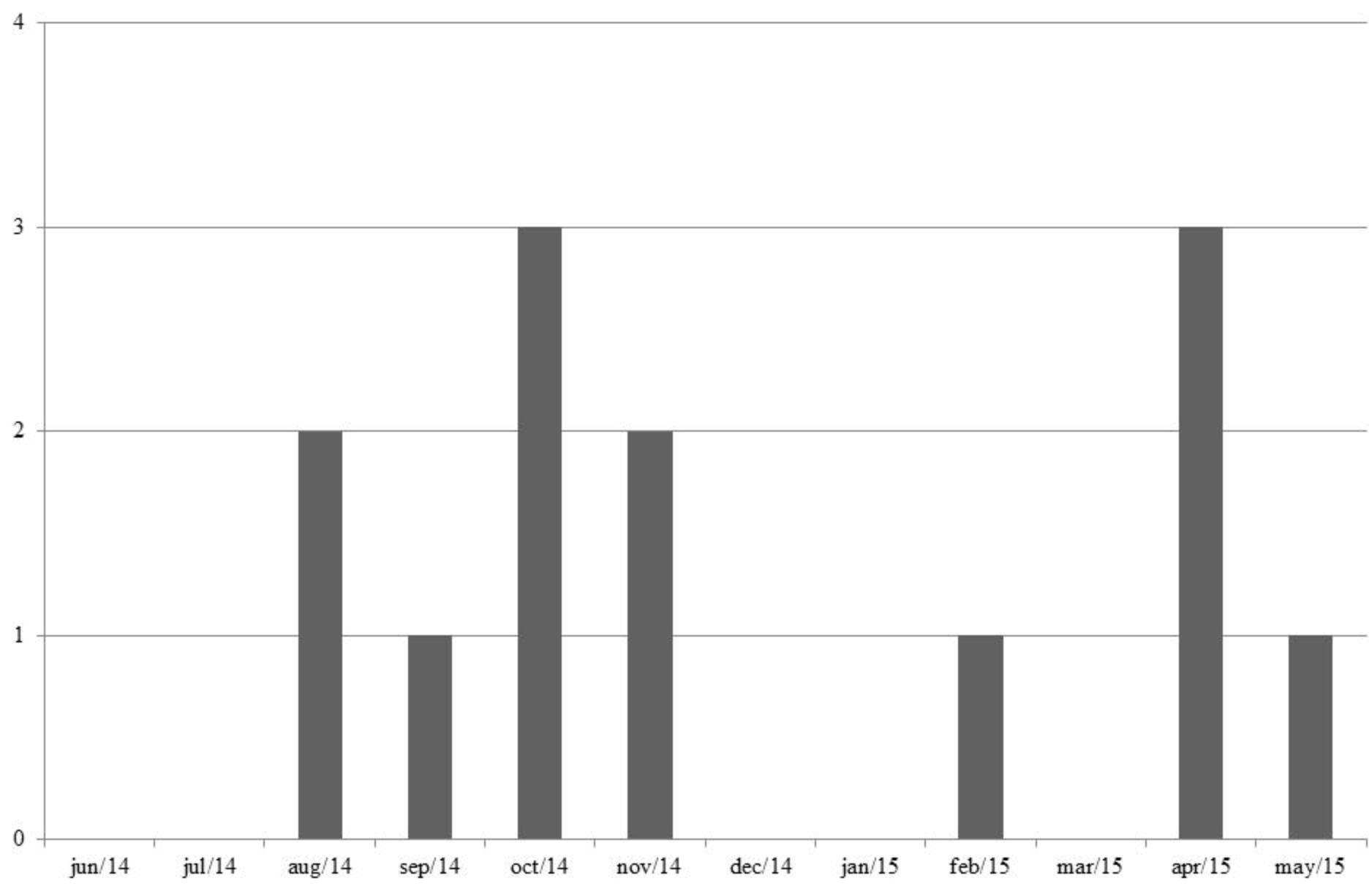


M: 50 bp molecular weight markers; **B1, B2, B3, B4, B5 and B6:** negative controls (blanks); **Pv**: positive controls for *Plasmodium vivax* (100 bp bands)


A. (N.) lanei
A. (N.) rangeli
A. (A.) mediopunctatus
A. (N.) parvus
A. (N.) argyritarsis
A. (N.) albitarsis
A. (N.) lutzii
A. (N.) galvaoi
A. (K.) homunculus
A. (N.) evansae
A. (N.) triannulatus
A. (N.) strodei
A. (K.) cruzii
0%
10%
20%
30%
40%
50%
60%
70%
80%
90%
100%
CDC CANOPY
CDC GROUND
PD
SHANNON


may/15
apr/15
mar/15
feb/15
jan/15
dec/14
nov/14
oct/14
sep/14
aug/14
jul/14
jun/14
0%
10%
20%
30%
40%
50%
60%
70%
80%
90%
100%
A. (K) cruzii
A. (N) strodei
A. (N) triannulatus
A. (N) evansae
A. (K) homunculus
A. (N) galvaoi
A. (N) lutzii
A. (N) albitarsis
A. (N) argyritarsis
A. (N) parvus
A. (A). mediopunctatus
A. (N) rangeli
A. (N) lanei


40,0
35,0
30,0
25,0
20,0
15,0
10,0
5,0
0,0
Temperature (°C)
250
200
150
100
50
0
Rainfall (mm)
jun/14
jul/14
aug/14
set/14
oct/14
nov/14
dec/14
jan/15
feb/15
mar/15
apr/15
may/15
Date (month/year)
Temperature (°C)
Rainfall (mm)


4
3
2
1
0
jun/14
jul/14
aug/14
sep/14
oct/14
nov/14
dec/14
jan/15
feb/15
mar/15
apr/15
may/15