

Malaria-transmitting *Anopheles* in the Colombian Pacific region

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INTRODUCTION

Malaria remains a significant public health concern in Colombia, and particularly, in the most malaria-endemic Pacific region¹. The detection of *Plasmodium* infected mosquitoes is an important parameter for the incrimination of vectors in malaria transmission². Therefore, the aim of this study was to detect natural infection by *Plasmodium* parasites in *Anopheles* mosquitoes collected in localities of the Colombian Pacific region.

METHODOLOGY

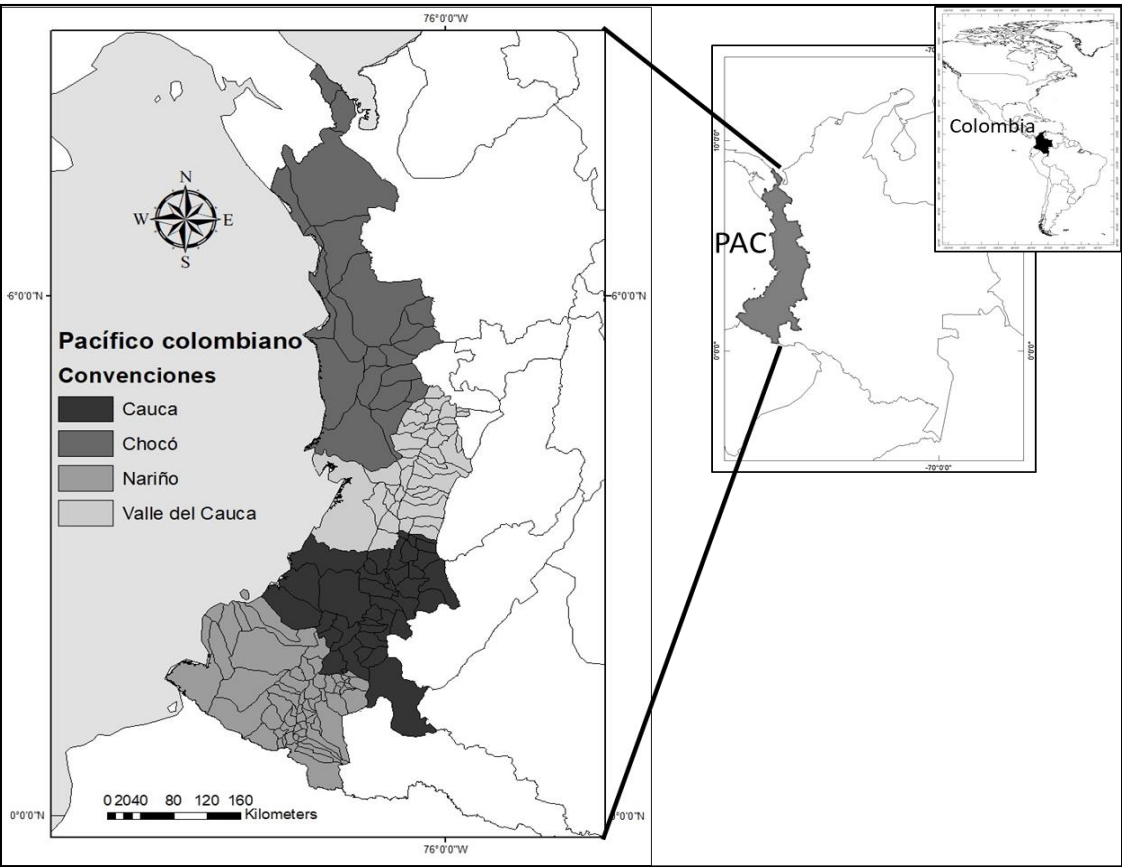


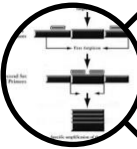
Figure 1. Pacific Region of Colombia. Sampled sites: San Antonio and Basurú, Istmina (Department of Chocó); Buchely and Inguapi del Carmen, Tumaco (Department of Nariño); La Playa and Salahondita, Francisco Pizarro (Department of Nariño); Córdoba and Zacarías, Buenaventura (Department of Valle del Cauca)



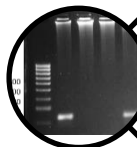
Collection of *Anopheles* mosquitoes intra and peridomicile from 18:00 - 24:00 hr



Morphological identification³. Molecular confirmation of species assignation (PCR-RFLP-ITS2) ⁴



Plasmodium detection by genus-specific nested PCR-18s ⁵

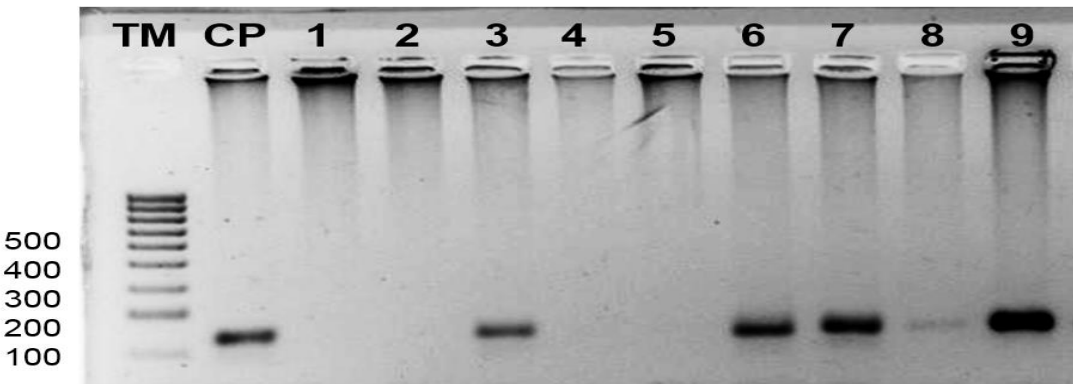


Plasmodium detection by genus-specific PCR-COXIII ⁶

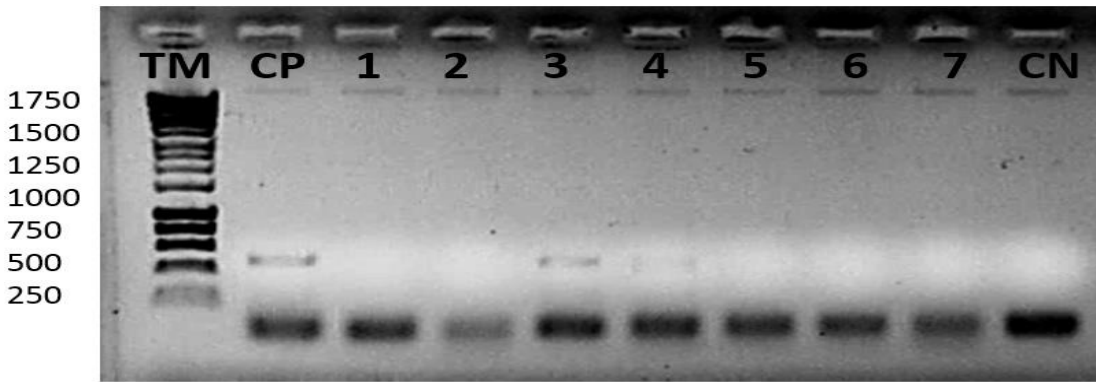
RESULTS

Table 1. Rate of *Anopheles* natural infection by *Plasmodium* spp.

Locality, Municipality (Department)	Species	n (%)	Parasite	Infection rate (%)
Córdoba, Buenaventura (Valle del Cauca)	<i>Anopheles nuneztovari</i>	715 (100%)	<i>Plasmodium</i> sp. ^a	0.14
Bucheley, Tumaco (Nariño)	<i>Anopheles albimanus</i>	191 (90.1%)	<i>P. falciparum</i> ^{a,b}	0.52
Salahondita, Francisco Pizarro (Nariño)	<i>Anopheles calderoni</i>	49 (40.2%)	<i>Plasmodium</i> sp. ^{a,b}	2.04
Basurú, Istmina (Chocó)	<i>Anopheles darlingi</i>	84 (47%)	<i>P. vivax</i> ^{a,b}	1.2



Nested PCR. CP: Positive control. **Samples 1-9.** Lane 6. *An. albimanus* positive for *P. falciparum* from Tumaco. Lane 8. *An. calderoni* positive for *Plasmodium* spp. Lane 9. *An. darlingi* positive for Istmina *P. vivax*. Electrophoresis, 0.8% agarose gel. TM: 1 Kb GeneRuler molecular marker



COXIII PCR. CP: Positive control. CN: Negative control. **Samples 1-7.** Lane 3. *An. albimanus* positive para *P. falciparum* from Tumaco. Lane 4. *An. darlingi* positive para *P. vivax* from Istmina. Electrophoresis, 0.8% agarose gel. TM: 1 Kb GeneRuler molecular marker

Figure 2. *Anopheles* natural infection by *Plasmodium* spp. using molecular methods.

CONCLUSIONS

The findings suggest that the main malaria vectors, *An. albimanus*, *An. darlingi* and *An. nuneztovari*, drive transmission in this region, but their distribution varies spatially.

The detection of *Plasmodium* natural infection in *An. calderoni* suggests its role as a local vector in specific areas.

These findings indicate that malaria transmission and vector roles vary spatially, emphasizing the need for locally adapted vector control interventions.

ACKNOWLEDGMENT

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