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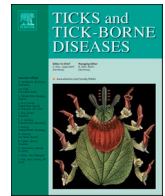
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Prevalence of heartwater in Guadeloupe (2024): stable endemicity and evidence of spread to Les Saintes

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ABSTRACT

Heartwater is a tick-borne disease caused by *Ehrlichia ruminantium* and remains endemic in the Guadeloupe archipelago. While previous data from 1989 suggested a seroprevalence of 22 % in cattle, no updated figures were available for livestock, and certain islands such as Les Saintes were considered disease-free.

This study aimed to update the sero-epidemiological status of heartwater in cattle and goats across Guadeloupe in 2024. Blood samples were collected from 261 cattle and 135 goats from all islands. Serological testing using the MAP1B ELISA was performed, and true seroprevalence was estimated using a simplified stochastic model that accounts for test sensitivity, specificity, and sample uncertainty. The mean cattle seroprevalence was estimated at 28 % (95 % CI [22–34 %]), with no significant differences between Basse-Terre, Grande-Terre, and Marie-Galante. Importantly, seropositive goats were identified on all major islands, including Les Saintes. Our findings suggest that heartwater remains endemically stable in Guadeloupe since the 1980s and reveal for the first time its serological presence in Les Saintes. These results emphasize the importance of strengthening regional surveillance systems, including veterinary reporting, serological monitoring, and coordinated efforts to prevent further spread across the Caribbean.

1. Introduction

Heartwater is a severe, often fatal, infectious disease affecting wild and domestic ruminants, caused by the intracellular bacterium *Ehrlichia ruminantium*, and transmitted by *Amblyomma* tick. It is endemic in sub-Saharan Africa, the Indian Ocean and Caribbean Islands, especially the Guadeloupe archipelago and Antigua. The tropical bont tick, *Amblyomma variegatum*, the principal vector of heartwater in the Caribbean, was introduced into the region in the 19th century and rapidly established across many islands (Barre and Uilenberg, 2010; Maillard et al., 1993). The first confirmed outbreaks of heartwater in the Caribbean were reported in the 80 s (Barré et al., 1984; Birnie et al.,

1985; Perreau et al., 1980; Uilenberg et al., 1984).

The presence of the vector and the pathogen in the Caribbean has raised concerns about the potential spread to other islands and the American mainland (Kasari et al., 2010). In response, eradication programs, such as the Caribbean *Amblyomma* Program (CAP) (Pegram et al., 1998), were initiated in the early 2000's. These efforts, based primarily on systematic acaricide treatment of animals, ultimately failed to eliminate *A. variegatum* from the Caribbean region due to various reasons (Ahoussou et al., 2010; Pegram et al., 2007). In 2008, studies conducted in Guadeloupe archipelago revealed a high prevalence of *E. ruminantium* in tick populations (Molia et al., 2008; Vachiéry et al., 2008). However, the last study assessing the seroprevalence of

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heartwater in livestock in Guadeloupe dates back to Camus (1989) and reported 22 % of heartwater seropositive cattle, though without specifying the diagnostic method used (Camus and Barre, 1987). Moreover, these studies described an endemic situation of the disease in Guadeloupe. Indeed, according to Deem et al. (1996), endemic stability is maintained by the convergence of three key factors: (1) high infection prevalence in the tick vector population, (2) the presence of reservoir hosts that sustain pathogen circulation, and (3) host infections occurring when host immunity is sufficiently high.

The present study aimed to provide an updated assessment of heartwater in the Guadeloupe archipelago in 2024 by estimating the seroprevalence in cattle and goats across all major islands, including previously non-endemic areas such as Les Saintes. Particular emphasis was placed on applying a robust statistical methodology to infer population-level prevalence while accounting for diagnostic test performance and sample uncertainty. Estimation of true seroprevalence at the population level from sampled data is a central task in veterinary epidemiology and requires inference modeling.

In this study, we applied a simplified stochastic approach based on Vose (2008) to estimate the distribution of the true prevalence among tested animals. This allowed to calculate the true number of infected animals in our sample. Using the same stochastic approach, we also estimated the population level seroprevalence, accounting for both the sample size and the imperfect test used, including the method used for its validation.

2. Methods

2.1. Study area

Guadeloupe, part of the lesser Antilles and a region of the French West Indies (FWI), is an archipelago comprising five islands: Basse-Terre and Grande-Terre islands constitute the “mainland” Guadeloupe in conjunction with three smaller islands, namely La Désirade, Marie-Galante and Les Saintes. The archipelago has a predominant tropical climate, characterized by two distinct micro-climates determined by important topographic differences between the two main islands. A rainier climate is observed in Basse-Terre mostly driven by trade winds and relief (Great Soufriere volcano massif, summiting at 1 467 m) while the Grande Terre and the other islands have a drier climate explained by absence of marked relief (Bleuse and Mandar, 1993). The livestock sector in Guadeloupe comprised around 34 800 cattle, 10 300 goat and 1 400 sheep in 2020 (Agregate, 2021). The traditional way to raise cattle in Guadeloupe is mainly tethered (Salas et al., 1986). However, this sector is declining due to various factors, including the threat caused by ticks and their associated diseases (Galan et al., 2008).

2.2. Sampling procedure

A complete list of 4 971 breeders (34,790 head of cattle) was obtained from local authorities (DAAF Guadeloupe: Direction of Alimentation, Agriculture and Forest). Considering the intra-herd transmission due to the shared tick-infested environment, a cluster sampling of animals was decided. Based on an expected seroprevalence of 20 % (Camus, 1989), and assuming a 5 % error ($\alpha=0.05$), an initial sample size of 246 cattle was calculated (Dohoo et al., 2014). A design effect of 1.15 was calculated based on an intra-cluster correlation coefficient of 0.05 for heartwater (Kenaw, 2023) and a cluster size corresponding to the median number of animals per breeders ($n = 4$). This adjustment resulted in a final required sample size of 283 cattle. To determine the numbers of farms where samples should be collected, the total number of animals was divided by the median herd size, yielding a target of 71 farms.

Sampling was conducted across the Guadeloupean archipelago from September 2023 to June 2024. The farms were sampled using a stratified sampling approach, with the proportion of farmers in each municipality serving as the sampling criterion (Dohoo et al., 2014). In some areas,

convenience blood sampling was performed for goats, specifically for the purposes of disease detection. This sampling was applied in regions where cattle were absent, such as the islands of Les Saintes and La Désirade. A total of 21 goat farms with 135 animals were sampled.

Animals were sampled after obtaining farmer’s consent. The study protocol received formal ethical approval prior to implementation (APAFIS #43,250–2023,050,210,511,531 v3). Blood samples were collected from restrained animals via either the jugular or the coccygeal vein, and transported under refrigerated conditions to the laboratory.

2.3. Serological survey in cattle and goats

Serum samples from cattle and goats were tested using a modified version of the reference ELISA test for the detection of antibodies specific to the *E. ruminantium* MAP-1B protein, a highly specific antigen previously described by Mondry et al. (1998) and Van Vliet et al. (1995). Briefly, MAP-1B antigen was coated on microplates. After blocking, sera and positive or negative controls were incubated. Plates were washed with PBS 0,05 % Tween20 and a HRP conjugate was added. After incubation and the last washings, TMB was added and specific colour developed if anti-MAP-1B antibodies were previously fixed. OD was measured at 450 nm and sera were declared positive if the Relative Positivity Percentage ($RPP = OD \text{ sample} / OD \text{ positive control} \times 100$) exceeded 10 % for goats and 24 % for cattle (Rodrigues, personal communication).

2.4. Statistical analysis

In epidemiology, estimating the true proportion at the population level based on a single sample relies on the assumption that the sample proportion follows a distribution. The true estimate of the mean from a single sample is calculated as $\hat{P} = r/n$, where r is the observed number of successes and n is the sample size. The standard variation of the true estimate of the mean is called the standard error, which is given by $[p(1-p)/n]^{1/2}$. Under specific conditions, a large sample size ($n > 30$), a prevalence within the range $0.05 \leq P \leq 0.95$, and both nP and $n(1-P) \geq 5$, the binomial distribution of the true estimate of the mean of the proportion can be approximated by a normal distribution (Thrusfield, 2007). This approach accounts for sampling uncertainty but is inaccurate for small sample size or for diseases with low prevalence. In addition, estimating the true prevalence becomes more complex when the diagnostic test used is not 100 % sensitive and specific (Rogan and Gladen, 1978). Abundant literature was published to determine confidence interval that adjust for imperfect diagnostic tests (Greiner and Gardner, 2000; Lang and Reiczigel, 2014; Reiczigel et al., 2010). The adjustment of the confidence interval of the true prevalence is obtained from the adjustment of the apparent prevalence and its confidence interval, as well as from the adjustment of the sensitivity and specificity when they are estimated from samples. In this deterministic approach, the adjustments systematically use the approximation of the normal distribution. More recently, Flor et al. (2020) compared these frequentist methods with a stochastic (Bayesian) approach for estimating prevalence.

Following serological testing, cattle sera were classified as either positive or negative, indicating recent infection within the last year. Doubtful results were considered negative to prioritize specificity, as it could indicate older infections (Semu et al., 2001). Differences in seroprevalence between the three main islands, Basse-Terre, Grande-Terre and Marie Galante, were assessed using the chi-square test (χ^2 -test). Statistical significance was defined for p-values < 0.05 .

An original method incorporating uncertainty was employed to estimate the prevalence of heartwater at population level. The apparent prevalence (AP) was first calculated by dividing the number of seropositive animals by the total number of animals tested (ns). The true prevalence (TP) was then estimated using the standard formula (Eq. (1)),

as described by Dohoo et al. (2014):

$$TP = \frac{AP + SP - 1}{SE + SP - 1} \quad (1)$$

With SP and SE being the specificity and sensitivity of the MAP-1B test. To consider their uncertainty, both specificity and sensitivity were modelled using beta distributions with uniform Beta (1,1) non informative prior distribution, with parameters $shape\ 1 = S + 1$ and $shape\ 2 = N - S + 1$ (Vose, 2008). In these equations, S represents the number of sick animals that tested positive (for sensitivity) or the number of healthy animals that tested negative (for specificity) and N denotes the total number of diseased animals tested (for sensitivity) or the total number of healthy animals tested (for specificity) during the test validation study. Validation data were obtained from the WOA reference laboratory for heartwater (Rodrigues, personal communication). The distribution of TP obtained was then multiplied by the number of animals tested (ns) resulting in a distribution of the true number of positive animals (nsp). To extrapolate to the entire Guadeloupean cattle population, the heartwater seroprevalence was modelled using a beta distribution with parameter $shape\ 1 = nsp + 1$ and $shape\ 2 = ns - nsp + 1$. From this distribution, the mean, 95 % confidence intervals, and skewness were calculated. This method was applied to ELISA results from cattle for the whole Guadeloupe and separately for the three major islands (Basse-Terre, Grande-Terre and Marie-Galante). A classical one-dimensional Monte-Carlo simulation with 10 000 iterations was performed for stochastic modelling.

Data were stored using Excel spreadsheets (Microsoft Corporation, 2021) and statistical analysis were conducted using R software (version 4.4.3.), with the `mc2d` package (version 0.2.1) (Pouillot and Delignette-Muler, 2010; R Core Team, 2023).

2.5. Geospatial interpolation

Geospatial interpolation was used to visualize the spatial distribution of heartwater across the Guadeloupe archipelago. A georeferenced layer of breeding locations was created, with each breeder represented by the centroid of their animals' geographical positions. Farms were classified as positive if at least one animal tested seropositive for heartwater. Data from 21 goat farms were integrated into the cattle farmers' dataset. To provide a spatial representation of heartwater seroprevalence across the Guadeloupean archipelago, Inverse Distance Weighting (IDW) interpolation was applied using a power parameter of 4 (Longley et al., 2015). This value was selected to interpolate the presence of heartwater within the Guadeloupean territory whilst ensuring the confidentiality of individual farm locations.

3. Results

Sample collection was conducted on 74 cattle farms and sera were collected from 261 cattle. *E. ruminantium* MAP-1B specific antibodies were detected in 15 out of 56 animals sampled in Basse-Terre, resulting in an apparent seroprevalence of 26.7 %. In Grande-Terre, 38 out of 137 animals tested positive, yielding an apparent seroprevalence of 27.7 %. In Marie-Galante, 23 of 68 animals were positives, resulting in an apparent seroprevalence of 33.8 %. Overall, the combined apparent seroprevalence across all sampled cattle was 29.1 % (76 out of the 261 tested cattle). Despite minor variations, the apparent seroprevalences across the three islands were comparable, and no statistically significant differences were observed between them, indicating a relatively uniform circulation of *E. ruminantium* in the cattle population across the archipelago (Table 1).

The distribution of cattle true seroprevalence modelled at population level presented a mean of 0.28 with a 95% confidence interval of [0.22; 0.34]_{95%} and a skewness of - 0.01 for the whole Guadeloupe, 0.27 [0.16; 0.39]_{95%} with a skewness of 0.26 for Basse-Terre, 0.27 [0.20; 0.36]_{95%} with a skewness of 0.16 for Grande-Terre and 0.34 [0.23; 0.45]_{95%} with

Table 1

Seroprevalence of *E. ruminantium* in cattle by island: results of the MAP-1B ELISA test performed on cattle sampled from the three major islands of the Guadeloupe archipelago. A total of 261 animals were tested: 56 from Basse-Terre, 137 from Grande-Terre, and 68 from Marie-Galante.

Island	ELISA +	ELISA -	%
Basse-Terre	15	41	26.7
Grande-Terre	38	99	27.7
Marie-Galante	23	45	33.8
Total	76	185	29.1

a skewness of 0.11 for Marie-Galante. Taken together, these results support the conclusion that heartwater seroprevalence remains consistently endemic across the Guadeloupe archipelago without marked geographical heterogeneity. (Figure 1).

Amongst the 74 cattle farms selected, 47 had at least one animal that tested positive for *E. ruminantium* using the MAP-1B serological test. The MAP-1B ELISA was also performed on 135 goats' serum samples collected from 21 farms located in Basse-Terre, Grande-Terre, Marie-Galante, la Désirade, and les Saintes. Amongst them, 27 seropositive animals were detected on all investigated islands, further supporting the broad geographic distribution of heartwater among ruminants in Guadeloupe.

The inverse distance weighting map further supported the widespread presence of heartwater, showing evidence of infection on farms across all islands within the archipelago (Fig. 2). Seropositive farms (goats or cattle) were detected across all major islands, including Basse-Terre, Grande-Terre, Marie-Galante, La Désirade, and, notably, Les Saintes. High seropositivity zones were especially concentrated in the central and northern parts of Grande-Terre, the southwestern and eastern parts of Marie-Galante, and several lowland regions of Basse-Terre (Fig. 2). Municipalities not included in the sampling are depicted in grey on the map, while the Guadeloupe National Park is highlighted in dark green. As expected, no samples were collected from within the National Park boundaries due to livestock absence in these protected areas. The spatial distribution of seropositive farms confirms the widespread circulation of *E. ruminantium* throughout the archipelago, with no major exclusion zones identified.

The violin plot displays the beta-distributed estimates of

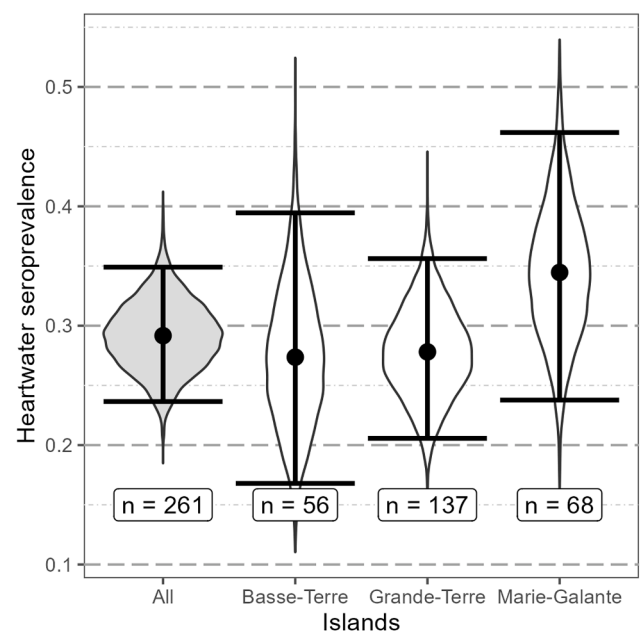


Fig. 1. Modelled heartwater seroprevalence in cattle across the Guadeloupe archipelago.

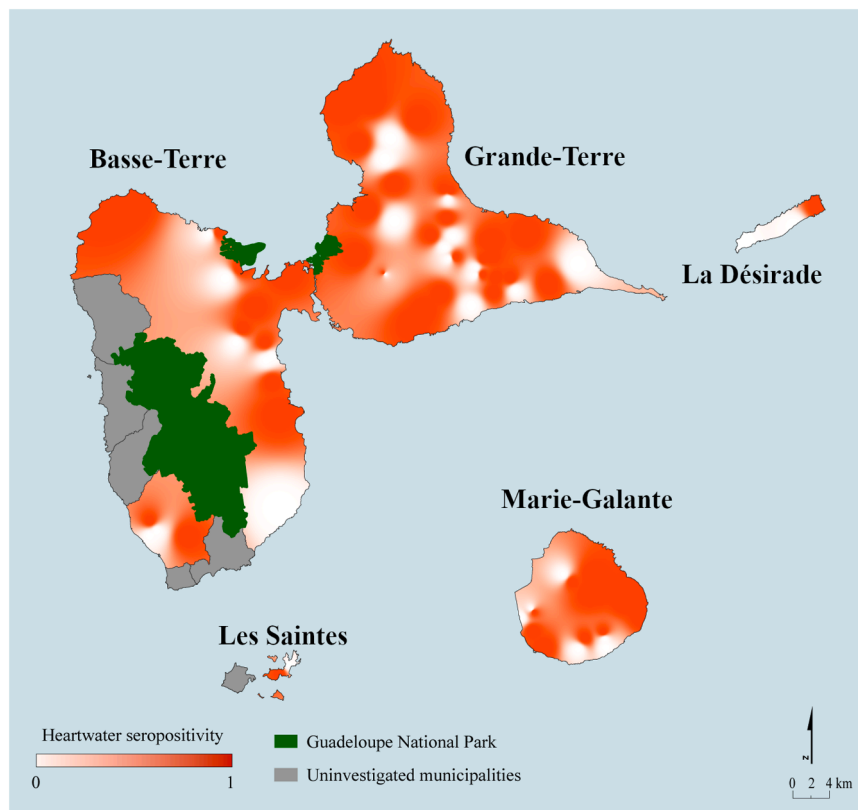


Fig. 2. Spatial distribution of heartwater seropositivity in cattle and goats across the Guadeloupe archipelago. This map presents the interpolated spatial distribution of heartwater seropositivity based on MAP-1B ELISA results from cattle and goat farms. Inverse Distance Weighting (IDW) interpolation was applied to farm-level data to visualize the relative intensity of seropositivity across the region. The color gradient ranges from white (no seropositivity, value = 0) to deep red (seropositivity evidence, value = 1).

E. ruminantium seroprevalence in cattle, modelled for the main islands of the Guadeloupe archipelago—Basse-Terre, Grande-Terre, and Marie-Galante—as well as the combined dataset ("All"). Each distribution was derived from stochastic modelling, incorporating diagnostic test uncertainty and sample size. Dots represent the mean seroprevalence, bars indicate the 95 % confidence interval, sample sizes (n) are indicated for each group. The density curves are depicted by the shapes around the centre lines.

4. Discussion

The results of this study revealed a heartwater seroprevalence of 0.28 (95 % CI [0.22; 0.34]) in cattle across the Guadeloupe archipelago. The simplified stochastic method to infer population-level seroprevalence produced a relatively symmetric distribution for the whole Guadeloupe, and slightly right-skewed distributions for individual islands. However, statistical analysis of the raw data did not reveal any significant differences in apparent seroprevalence between Basse-Terre, Grande-Terre, and Marie-Galante. These results are consistent with the 22 % seroprevalence reported by Camus in 1989 (lower limit of the current confidence interval) reinforcing the hypothesis of epidemiological stability of *E. ruminantium* circulation in Guadeloupe over the past 35 years.

The consistent circulation of *E. ruminantium* in Guadeloupean cattle suggests that heartwater is endemically stable, potentially maintained by repeated exposure. Similar results have been reported among cattle in endemic regions of West and Southern Africa, where seroprevalence detected using the MAP-1B ELISA test range from 20 % to 40 % (Bell-Sakyi et al., 2003; Mahan et al., 1998; Peter et al., 2002). The transient nature of the antibody response to the MAP-1B antigen, especially under conditions of recurrent natural exposures, may result in moderate but stable seroprevalence levels over time (Semu et al., 2001).

Therefore, these data provide strong evidence that *E. ruminantium* was actively circulating in the Guadeloupean cattle population during 2023–2024.

Serological evidence of heartwater was also detected in goats across all surveyed islands, including La Désirade and Les Saintes. The detection of seropositive animals in Les Saintes, a region historically considered free of heartwater, suggests a recent introduction and ongoing spread of the pathogen beyond previously recognized endemic areas. Based on veterinary field investigations and animal movement records, the most plausible hypothesis is the introduction in 2015 of infected livestock or animals infested with infected ticks from Marie-Galante. At the end of 2024, nervous clinical signs on goats were reported by farmers and the mandated veterinarian. These cases were subsequently confirmed by molecular diagnosis (unpublished data) which supports the active presence of *E. ruminantium* in Les Saintes. These findings highlight the successful establishment and active dissemination of the disease across all major islands of Guadeloupe. While many other Caribbean islands are currently infested with *A. variegatum* ticks (Charles et al., 2020; Kasari et al., 2010), most remain free of heartwater, as evidenced by the absence of reported clinical signs or molecular detection of *E. ruminantium*. The last study on sheep, cattle, and goats from various Caribbean islands, conducted in 2011, reported <1 % seroprevalence with no clinical cases, but these findings were then attributed to false positive reactions probably due to cross-reactivity of MAP-1B antigen with other tick-borne ehrlichiosis (Kelly et al., 2011). Nonetheless, this assertion may be subject to scrutiny in view of the supporting references to earlier studies (Kelly et al., 1994; Muller Kubold et al., 1992; Savadye et al., 1998) which used different methodologies to detect *E. ruminantium* antibodies. The validation paper for the MAP-1B ELISA (Mondry et al., 1998) clearly showed a significantly improved specificity of this test compared to crude or MAP-1 antigens

tests.

The recent detection of seropositive goats in Les Saintes, an area previously considered free from heartwater, underscores the need to maintain targeted surveillance in the region. Establishing a robust syndromic surveillance system, involving veterinary practitioners, is critical for the early detection of heartwater outbreaks and for tracking seasonal or annual variations in disease prevalence (Thompson and Etter, 2015). By systematically recording clinical cases, tick infestations, suspected heartwater occurrences, seroprevalence and *E. ruminantium* detection in ticks, veterinary services could provide valuable epidemiological data to inform both preventive and control strategies. In Guadeloupe, such a program was implemented with the RESPANG (REseau de Surveillance des Pathologies Nerveuses chez les ruminants de Guadeloupe) project in 2010 (Vachière et al., 2011). An innovative surveillance network was established to collect and diagnose ruminants showing nervous syndromes. This network involved veterinary practitioners, research sector and local authorities (Lefrançois et al., 2011), but unfortunately, RESPANG is not active anymore.

The spread of heartwater in Guadeloupe archipelago highlighted the necessary cooperation between Caribbean islands. This collaboration should focus on understanding the introduction dynamics and tracking the spread of infected ticks and/or livestock between islands. Potential risk factors analysis would improve recommendations to control heartwater and mitigate tick infestations. A coordinated effort involving local farmers, veterinary services, and epidemiologists will be essential to prevent further spread of the disease to neighbouring islands. Such cooperation already existed in the Caribbean with the Caribbean Amblyomma Program (CAP) (Barré et al., 1996; Pegram et al., 1998) which aimed to eradicate *A. variegatum* from the Caribbean but failed due to numerous factors (Ahoussou et al., 2010; Pegram et al., 2004, 2007). Today, the *CaribVET*¹ network is ideally positioned to coordinate and facilitate such initiatives. Leveraging its established connections with veterinary authorities and research institutions across the Caribbean, *CaribVET* could promote standardized protocols for sampling, diagnosis, and data sharing. This coordinated approach would strengthen the region's ability to early detect outbreaks, assess epidemiological trends, and implement effective control measures. Despite existing vector control measures promoted by the Guadeloupean sanitary defence group (SANIGWA), the persistence of *A. variegatum* populations and the continued spread of heartwater underscore the need for more effective tick management strategies. Moreover, accelerating the development of a heartwater vaccine remains a key priority if long-term control of this economically significant disease is to be achieved.

5. Conclusion

This study confirmed that heartwater remains endemic in Guadeloupe. Those results represent the first estimation of Guadeloupean cattle seroprevalence using the WOA reference ELISA. An original statistical method was applied to determine heartwater prevalence at the cattle population level. Stable levels of seroprevalence have been observed on the main islands. Furthermore, there is evidence to suggest that the rickettsial agent have spread to previously unaffected areas, such as Les Saintes. These findings underscore the need for strengthened regional surveillance, control strategies and partnership between actors of the livestock sector. Reactivation of local network, such as the previously existing RESPANG, appears determinant to efficiently collect valuable data on disease's epidemiology.

Given the continued presence of *A. variegatum* across the Caribbean and the risk of further geographic expansion, coordinated efforts in tick management and disease prevention at a regional scale are essential. Investment in research, including tick control, vaccine development and regional monitoring, will be critical to safeguarding livestock health and

maintaining agricultural resilience across the Caribbean and beyond. By combining local and regional surveillance, stakeholders can better anticipate epidemiological trends, limit disease spread and safeguard livestock industries throughout the Caribbean region.

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CRedit authorship contribution statement

Victor Dufleit: Data curation, Formal analysis, Investigation, Methodology, Software, Validation, Visualization, Writing – original draft. **Laure Guerrini:** Writing – review & editing, Visualization, Methodology, Formal analysis. **Mélanie Dhune:** Resources, Methodology, Investigation. **Laetitia Viry:** Resources, Methodology, Investigation. **Armande Belfort:** Resources, Methodology, Investigation. **Loïc Jacquet-Crérides:** Investigation. **Jimmy Duddy:** Investigation. **Joanna Cozier:** Writing – review & editing, Investigation, Data curation. **Michel Naves:** Supervision, Methodology, Investigation, Funding acquisition, Conceptualization. **Ludovic Arthein:** Investigation. **Thony Samut:** Investigation, Funding acquisition, Conceptualization. **Sylvie Lecolinet:** Methodology, Validation, Supervision. **Manuel Pezeron:** Investigation, Funding acquisition, Conceptualization. **Valerie Rodrigues:** Writing – review & editing, Writing – original draft, Supervision, Methodology, Conceptualization. **Damien F Meyer:** Conceptualization, Methodology, Supervision, Writing – original draft, Writing – review & editing. **Eric M C Etter:** Writing – review & editing, Writing – original draft, Validation, Supervision, Project administration, Methodology, Conceptualization.

Declaration of competing interest

The authors declare that they have no financial conflicts of interest in relation to the content of the article.

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Data availability

The data that has been used is confidential.

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