Risk factors associated with the frequency of antibodies against *Babesia bovis* and *Babesia bigemina* in cattle in southern Mozambique

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The study aimed to evaluate the risk factors associated with the frequency of IgG antibodies against *Babesia bovis* and *B. bigemina* in cattle in southern Mozambique. Eight hundred and nine serum samples were collected from cattle in three provinces namely Maputo, Gaza and Inhambane, and tested by indirect enzyme-linked immunosorbent assay (i-ELISA) to assess the humoral immune response towards *B. bovis* and *B. bigemina*. The chi-square test at 5% significance was used to determine whether there was an association between gender, age and geographic origin of seropositive animals. The overall prevalence was 78.8% (548/695) for *B. bovis* and 76.0% (528/695) for *B. bigemina*. The origin of the animals showed a significant association (p<0.05) with seropositivity to both agents, while gender and age was not associated (p>0.05). Maputo province had the highest rate of positive animals, with 93.7% (118/126) for *B. bovis* and 97.6% (123/126) for *B. bigemina*. In Gaza province 77.3% (321/415) of the animals were positive for *B. bovis* and 67.5% (280/415) for *B. bigemina* while in the province of Inhambane the levels of seropositivity were 70.8% (109/154) and 81.2% (125/154) for *B. bovis* and *B. bigemina* respectively. In the present study, the frequency of cattle positive for *B. bovis* and *B. bigemina* was shown to increase among older age groups, suggesting that infection and re-infection persisted even after the primary infection. Thus, this region is considered to be in a state of enzootic stability with regards to *B. bovis* and *B. bigemina*.

INDEX TERMS: *Babesia bovis*, *Babesia bigemina*, cattle, babesiosis, epidemiology, Mozambique.
**INTRODUCTION**

About 902,579 cattle are reared in the southern region of Mozambique, a value equivalent to 53.6% of the national total, estimated at 1,683,589 animals (Tia, 2008). These animals are prone to infection by *Babesia bovis* and *B. bigemina*, which are directly associated with the presence of the ticks *Rhipicephalus microplus*, *R. decoloratus*, *R. erson ets everts* and *Hyalomma marginatum rufipes* (De Matos 2008).

In the mid-1990s the government of Mozambique implemented a livestock restocking program based on the import of cattle from the Republic of South Africa and Zimbabwe (Martins et al. 2008). However, this increase in the production of ruminants in the country was not accompanied by improvements in sanitary conditions, which led to the occurrence of different diseases including the following tick-borne conditions: Babesiosis, anaplasmosis, ehrlichiosis and theileriosis. These diseases negatively affected the productivity and reproductive efficiency of livestock by causing the death of more than half the animal population (Simuunza et al. 2010).

These pathogens trigger a massive destruction of red blood cells in their hosts, promoting lysis of these cells followed by invasion of other erythrocytes (De Vos et al. 2004). Thus, infections with *Babesia* spp. in cattle are characterized by fever, anemia, hemoglobinemia, hemoglobinuria, and in many cases, death (Martins et al. 2008).

Babesiosis is responsible for severe damage to the animal and exerts a large economic impact on livestock production in tropical regions (Jonsson et al. 2008). The dynamics of infection by *Babesia* spp. depend on several factors, such as size and composition of the population of ticks, the transmission capacity of the vector and the susceptibility of the cattle, which can vary with breed, age and physiological conditions (Bock et al. 1997, Jonsson et al. 2008) as well as the agro-ecological and edaphic-climatic conditions of each geographical region.

The seroepidemiological study of bovine babesiosis in a given geographical area is important, because it may reveal the potential for outbreaks of this disease. Serological diagnostic methods are essential tools for assessing the prevalence and the immune status of animals as well as generating important information that can be used to control the tick vectors (Mahoney & Ross 1972, Mahoney 1975). The indirect enzyme-linked immunosorbent assay (i-ELISA) is the test that has given better results due to its high level of sensitivity and specificity (Madruga et al. 2001).

The objective of the current investigation was to examine possible correlation between gender, age and geographic region as risk factors associated with the prevalence of IgG antibodies against *B. bovis* and *B. bigemina* in cattle from the provinces of Maputo, Gaza and Inhambane, southern Mozambique.

**MATERIALS AND METHODS**

This study was carried out in the southern region of Mozambique, which occupies an area of 167,641 km², at a latitude of 25°58” south and a longitude of 32°35’ east, with an average altitude of 120 meters, characterized by a tropical, humid climate (Inam 2010). The current evaluated cattle (*Bos indicus*) and their crossbreed from herds in the provinces of Maputo, Gaza and Inhambane. These provinces account for 53.6% of the national cattle population, equivalent to 902,579 head, with 179,028 in inhambane, 569,404 in Gaza and 154,147 in Maputo respectively (Tia 2008).

A total of 809 serum samples were collected. The samples were derived from 110 rural smallholders with 4 located in Maputo, 78 in Gaza and the remaining 28 in Inhambane. The study was carried out between January to March 2010, and cattle owners participated voluntarily in this study. The inference was performed by non-probabilistic sampling, obeying the proportional stratification of each province studied. Thus, in each rural smallholder community samples were taken in order to represent 10% of the total herd.

In order to assess the association between rates of IgG antibodies against *Babesia bovis* and *B. bigemina* and the variables of age, gender and origin, animals were categorized as follows: Age (≥ 6 to ≤ 12 months, and >12 to ≤36 months and > 36 months), sex (male and female) and origin (Maputo, Gaza and Inhambane).

Blood samples were collected aseptically by coccygeal venipuncture in 10mL tubes *vacutainer*. Subsequently, the samples from the province of Maputo were transported under refrigeration using ice packs to the Central Veterinary Laboratory (CVL), Directorate of Animal Science (DCA) at the Institute for Agrarian Research of Mozambique (IIAM), samples from Gaza were processed at the Regional Veterinary Laboratory of South-Gaza/IIAM while the samples from Inhambane were processed at the Provincial Livestock Services Laboratory of Inhambane. In all cases, samples were submitted to centrifugation at 2,000x-g for 10 minutes in order to separate the clot and obtain serum. About 1mL of serum was separated into polypropylene microtubes of 1.5ml and placed in a freezer at -20°C until the execution of serological tests. Serological analyses were performed at the Laboratory of Parasitic Disease, Department of Epidemiology and Public Health, Federal Rural University of Rio de Janeiro (UFRF) and at the Laboratory of the National Center for Research in Agrobiology, Brazilian Agricultural Research Corporation (Embrapa Agrobiologia). Serum samples were tested by i-ELISA for the detection of IgG antibodies against *B. bovis* and *B. bigemina* using 96 well microplate 96 (Corning, Costar® 3590, USA). The microplates were previously sensitized with crude antigen of *B. bovis* and *B. bigemina* kindly supplied by National Research Center for Beef Cattle of the Brazilian Agricultural Research Corporation (Embrapa Gado de Corte), and processed according to the technique described by Madruga et al. (2001). Plates were read using a microplate spectrophotometer (LABSYSTEMS IEMS Reader MF) at a wavelength of 492hm. The cutoff for i-ELISA was established with the confidence level of 99.5% Frey et al. (1998).

The frequency of seropositive cattle of all categories (age, gender and origin) were calculated and differences between groups assessed by using Chi-square ($\chi^2$) at 5% of significance using the statistical software *BioStat* ©, version 4.0 (Ayres et al. 2000).
and Gaza (p<0.05). Babesia spp. were more likely to be seropositive for B. bovis, whereas in the Maputo, Inhambane and Gaza the frequency (p<0.05) of seropositive cattle for both agents, significant differences between provinces showed the highest prevalence was 93.7% (118/126), 70.8% (109/154) and 77.3% (321/415) for B. bovis, and 97.6% (123/126), 81.2% (125/154) and 67.5% (280/415) for B. bigemina, respectively. The prevalence rate showed that cattle in the province of Maputo were more likely to be seropositive for B. bigemina and B. bovis than animals coming from the provinces of Inhambane and Gaza (p<0.05).

### RESULTS

Prevalence values of animals seropositive for Babesia bovis and B. bigemina in three provinces of southern Mozambique (Maputo, Gaza and Inhambane) are provided in (Table 1). The results show that the overall prevalences of antibodies against B. bigemina and B. bovis were 76.0% (528/695) and 78.8% (548/695) respectively.

The percentages of prevalence for Babesia spp. determined on the basis of age, gender and geographic region are represented in (Table 1). There was no association (p>0.05) between gender and age seropositivity for either B. bovis or B. bigemina, on the other hand, while the factors of origin of the animals showed a significant association (p<0.05) with seropositivity to both agents studied.

Analysis of data for B. bovis revealed a prevalence of 82.2% (97/118), 78.3% (90/115) and 78.1% (361/462) among animals of age ≥6 to ≤12 months, >12 to ≤36 and over 36 months, respectively, (Table 1). For B. bigemina the seroprevalence was 80.5% (95/118), 78.3% (90/115) and 74.2% (343/462) for the three age groups examined. However, no significant differences were observed among groups aged greater than 12 months p>0.05 (Table 1).

When evaluating the demographic region as a risk factor associated with seropositivity, we found the statistically significant differences between provinces showed the highest frequency (p<0.05) of seropositive cattle for both agents, whereas in the Maputo, Inhambane and Gaza the frequency was 93.7% (118/126), 70.8% (109/154) and 77.3% (321/415) for B. bovis, and 97.6% (123/126), 81.2% (125/154) and 67.5% (280/415) for B. bigemina, respectively. The prevalence rate showed that cattle in the province of Maputo were more likely to be seropositive for B. bigemina and B. bovis than animals coming from the provinces of Inhambane and Gaza (p<0.05).

### DISCUSSION

The results of this study concur with those of previous other conducted in countries from tropical regions (Jonsson et al. 2008). In regions in eastern and southern Africa, studies indicate that bovine babesiosis presents seropositivity ranging from 19.5 to 94.0% of the cattle (Dreyer et al. 1998, Mbata et al. 2002, Mtshali et al. 2004, Martins et al. 2008). In South Africa, these studies showed that the prevalence of antibodies against Babesia bigemina ranged from 62.4 to 94.0%, and in the case of B. bovis the frequency was 19.5%. In Kenya the prevalence of Babesia spp. in cattle was 37.1%, according to the study of Okuthe & Buyu (2006).

In southern Mozambique is raised predominantly beef cattle, consisting mostly of Indian origin breed (Bos indicus) and their crossbreed, which are considered to be naturally resistant to parasitism by ticks (Mahoney 1975). However, the prevalence in this region for B. bovis and B. bigemina was found to be 78.8% (548/695) and 76.0% (528/695) respectively, suggesting great potential for transmission of these agents by ticks prevalent in that area. According to Mahoney & Ross (1972), prevalence rates equal to or greater than 75.0% classify the area as one of enzootic stability. Furthermore, it indicates that the cattle are infected before completing the first year of life, and are constantly re-infected. In this way, the animals maintain their immune system active against the agents in question.

In present study the seroprevalence detected was higher than 75%, characterizing the southern region of Mozambique as an area of enzootic stability for bovine babesiosis. In a situation of enzootic stability, calves are infected during the early months of life and are protected by colostral antibodies (passive immunity), thus enabling the development of active immunity without the manifestation of clinical disease. However, it should be noted that a study examining calves with age below six months (Jonsson et al. 2008), it was concluded that under conditions of high tick infestation high morbidity and mortality can occur even in during the period when protection was being conferred by colostral antibodies (Jonsson et al. 2008).

The association of gender with the seropositivity of the cattle to B. bigemina and B. bovis was not significantly different (p>0.05), confirming the findings of previous studies (Soares et al. 2000, Trindade et al. 2010).

The statistically significant differences observed in positive cattle from the same provinces studied in relation to the prevalence of antibodies against B. bovis and B. bigemina, may be due to variation in the rate of infection by ticks and, subsequently, a variation in the rate of cattle inoculation by the infected ticks. The animals are exposed to more risk conditions associated with outbreaks of babesiosis infection by B. bovis and B. bigemina in cattle was not homogeneous among the same province.

In the current study, the seropositivity for B. bigemina 77.3% (321/415) in Gaza province was higher when compared with the level obtained for B. bovis 67.5% (280/415). In this province, there exists a higher percentage of Bos indicus species, which are considered to more resistant to tick infestation (Madruga et al. 1984), and with which higher larval mortality is observed (Gonzáles 1974).

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**Table 1. Prevalence of antibodies against Babesia bigemina and Babesia bovis in cattle detected by indirect enzyme-linked immunosorbent reaction and associated factors in Southern of Mozambique**

<table>
<thead>
<tr>
<th>Factor</th>
<th>N</th>
<th>Babesia bigemina</th>
<th>Babesia bovis</th>
<th>B. bigemina e B. bovis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>%</td>
<td>n</td>
<td>%</td>
</tr>
<tr>
<td>Prevalence</td>
<td>695</td>
<td>528</td>
<td>76.0</td>
<td>548</td>
</tr>
<tr>
<td>Age (months)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≥6 the ≤12</td>
<td>118</td>
<td>95</td>
<td>80.5&lt;sup&gt;a&lt;/sup&gt;</td>
<td>97</td>
</tr>
<tr>
<td>&gt;12 the ≤36</td>
<td>115</td>
<td>90</td>
<td>78.3&lt;sup&gt;b&lt;/sup&gt;</td>
<td>90</td>
</tr>
<tr>
<td>&gt;36</td>
<td>462</td>
<td>343</td>
<td>74.2&lt;sup&gt;a&lt;/sup&gt;</td>
<td>361</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Females</td>
<td>438</td>
<td>334</td>
<td>76.3&lt;sup&gt;a&lt;/sup&gt;</td>
<td>345</td>
</tr>
<tr>
<td>Males</td>
<td>257</td>
<td>194</td>
<td>75.5&lt;sup&gt;a&lt;/sup&gt;</td>
<td>203</td>
</tr>
<tr>
<td>Province</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Maputo</td>
<td>126</td>
<td>123</td>
<td>97.6&lt;sup&gt;a&lt;/sup&gt;</td>
<td>118</td>
</tr>
<tr>
<td>Inhambane</td>
<td>154</td>
<td>125</td>
<td>81.2&lt;sup&gt;b&lt;/sup&gt;</td>
<td>109</td>
</tr>
<tr>
<td>Gaza</td>
<td>415</td>
<td>280</td>
<td>67.5&lt;sup&gt;c&lt;/sup&gt;</td>
<td>321</td>
</tr>
</tbody>
</table>

N = numbers of animals collected, n = numbers of samples positive to Babesia bigemina and B. bovis. Values in the column followed by same letters do not have statistically significant difference between groups or factors studied (a) by test χ² (p<0.05). Values with statistically significant differences between provinces are indicated by different capital letters (A, B and C) by test χ² (p<0.05).
These data demonstrate that the disease has a varied distribution within the provinces. A previous study of natural infection in bovines of the beef cattle breed from the region the Tete province revealed a seropositivity level of 39.1% for *B. bovis* (Alfredo et al. 2005). It is possible that the observed differences between the earlier study and our data may have resulted from a seasonal increase in the tick population, in a lack of control of such an increase, or from the subsequent introduction of susceptible animals, derived from South Africa and Zimbabwe, during the livestock restocking program.

During the blood collection process, the presence of ticks was observed on all the farms, despite the frequent use of tick control drugs containing pyrethroids and avermectins, as well as a high prevalence of babesiosis. Host and environment are factors that affect the prevalence of babesiosis in a particular region (Mahoney & Ross 1972).

Information in relation to the levels of antibodies against *B. bovis* and *B. bigemina* in cattle in the study areas will provide insights to managers of national livestock to improve their knowledge of the immunological status of herds, and may contribute towards the development of future interventions and management strategies in animal health. In this context, knowing the prevalence of agents in each province, one can develop geo-referenced maps of regions of stability and instability to provide better control of enzootic disease in the country.

The results of this study corroborate the findings from studies conducted in the countries of southern and eastern Africa, where the variation of infection by *B. bigemina* and *B. bovis* depends on factors such as the prevalence of vectors and measures related to their control, but is independent of gender.

We conclude that the southern region of Mozambique should be considered as an area of enzootic stability for both *B. bovis* and *B. bigemina*. Moreover, this region offers risk for transmission of babesiosis to susceptible animals coming from areas of enzootic instability. Therefore, the use of appropriate preventive measures is needed, especially with regard to the effective control of ticks. In these circumstances, further studies would be justified in order to monitor the herds for purpose of identifying factors that may pose risks to the current epidemiological status of the region.

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