

Molecular-biological and serological tools for the diagnosis of canine and equine babesiosis in Switzerland

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Key words

Babesia, Theileria, Rhipicephalus, Dermacentor, PCR

Aim of the study

Babesiosis in dogs and horses is caused by protozoan parasites including *Babesia canis* for dogs, while horses are affected by *B. caballi* and *Theileria equi*. For a long time both, canine and equine babesiosis, were assumed to be typical travelers diseases, but within the last years more and more cases were found in patients that had no anamnestic sojourn outside Switzerland. Therefore, these parasites become more and more relevant for Swiss veterinarians and the local animal population. The aim of the present project was to prepare laboratory tools suitable to diagnose canine and equine babesiosis/prioplasmosis, and to apply those tools in an explorative study on (a) the occurrence of autochthonous and imported cases of babesiosis in dogs and horses in our country and (b) respective tick hosts.

Material and methods

For routine diagnosis of *Babesia*-infection in dogs and horses, investigation of Giemsa-stained blood-smears (capillary blood) for acute stage disease, and antibody detection by IFAT for chronic stage disease are the appropriate tools. Speciation of *Babesia* was performed by PCR, which was subsequently developed into a new real-time PCR, suitable for routine diagnosis, but also for testing Giemsa-stained blood smears and ticks.

Results and significance

Overall, 99 horses of Switzerland (either with clinically suspected babesiosis or at risk of infection) were tested in the present study. One of these horses yielded a positive Giemsa-stained blood smear at diagnosis, subsequently confirmed as *T. equi* by PCR. From 14 clinically suspected cases of canine babesiosis, 5 were parasitologically positive, and all 5 cases yielded *B. canis canis* by PCR.

A total of 1'864 ticks were collected from mainly dogs and horses. 34 (1.8%) specimen were *Dermacentor reticulatus*, 64 (3.4%) were *Rhipicephalus sanguineus* and the rest (94.8%) *Ixodes* spp.. All *R. sanguineus* ticks were negative by *Babesia*-PCR, while 3 (9%) out of the 34 *D. reticulatus* specimen were PCR-positive for *Babesia*-DNA. Subsequent sequencing revealed a high similarity with *Babesia* sp. Genotype EU1.

Global warming, ecological changes in the potential habitat of ticks, increasing host- and vector-populations and increasing mobility of dog owners may be responsible for an autochthonous risk of infection for *Babesia* spp., and further surveillance is suggested to tackle this problem.

Publications, posters and presentations

Porchet, MJ.; Sager, H.; Muggli, L.; Oppliger, A.; Müller, N.; Frey, C.; Gottstein, B. Etude épidémiologique descriptive de la Babésiose canine dans la Région Lémanique. Schweiz. Arch. Tierheilkd. 149: 457-465 (2007).

Porchet, MJ.; Sager, H.; Muggli, L.; Oppliger, A.; Müller, N.; Frey, C.; Gottstein, B. A descriptive epidemiological study on canine babesiosis in the Lake Geneva region. WAAVP, Gent/Belgium, August 19-23, 2007.

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