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Abstract: BACKGROUND: Angiostrongylus vasorum is a potentially fatal canine nematode. Due to the high variability of clinical signs and the often chronic and subtle course of the infections, the diagnosis is particularly challenging. A rapid in-clinic assay (Angio Detect™ Test, IDEXX Laboratories, Westbrook, Maine, USA) for the serological detection of circulating antigen and intended for routine in-clinic diagnosis has been evaluated. METHODS: Sensitivity was calculated with sera from 39 naturally infected dogs confirmed by Baermann-Wetzel analysis, while sera of 38 experimentally infected dogs were used for follow-up analyses, of which 10 were treated with imidacloprid/moxidectin. Cross-reactivity was tested with a total of 123 samples from dogs with proven parasitic infections with Toxocara canis (n = 21), Ancylostoma caninum (n = 4), Crenosoma vulpis (n = 18), Oslerus osleri (n = 3), Eucoleus aerophilus, (n = 6), Dirofilaria immitis (n = 28), Dirofilaria repens (n = 20), Acantocheilonema reconditum (n = 10) or Dipetalonema dracunculoides (n = 10) or multiple infections (n = 3). All sera were tested with the Angio Detect™ Test and with an ELISA for detection of circulating antigen of A. vasorum. RESULTS: The sensitivity of the Angio Detect™ Test was 84.6% (95% C.I. 69.5 - 94.1%), while specificity was 100% (95% C.I. 97.6 - 100%). The sensitivity of the ELISA (94.9%, 95% C.I. 82.7 - 99.3%) was comparable with previous evaluations. In experimentally infected dogs, earliest positive results with the Angio Detect™ Test were observed 9 weeks post inoculation and 5 weeks later all sera were Angio Detect™ Test positive. After anthelmintic treatment, seropositive dogs turned negative again within 3 to 7 weeks after treatment. The evaluation of the colour intensity of the test strips confirmed the delay of approximately 3-4 weeks for antigen detection by the Angio Detect™ Test compared to the ELISA and its correlation with the time after infection. CONCLUSIONS: This study provided evidence of a good sensitivity and a very high specificity of the rapid device Angio Detect™ Test for detection of circulating A. vasorum antigen in dogs with suspected canine angiostrongylosis, representing a very simple and useful tool to be broadly applied in veterinary practices. The rapid detection of infected dogs is a key point for initiating an indispensable and urgent therapy.

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Abstract
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Methods: Sensitivity was calculated with sera from 39 naturally infected dogs confirmed by Baermann-Wetzel analysis, while sera of 38 experimentally infected dogs were used for follow-up analyses, of which 10 were treated with imidaclorpid/moxidectin. Cross-reactivity was tested with a total of 123 samples from dogs with proven parasitic infections with Toxocara canis (n = 21), Ancylostoma caninum (n = 4), Crenosoma vulpis (n = 18), Oselus osleri (n = 3), Eucoleus aerophilus, (n = 6), Dirofilaria immitis (n = 28), Dirofilaria repens (n = 20), Acanthocheilonema reconditum (n = 10) or Dipetalonema dracunculoides (n = 10) or multiple infections (n = 3). All sera were tested with the Angio Detect™ Test and with an ELISA for detection of circulating antigen of A. vasorum.

Results: The sensitivity of the Angio Detect™ Test was 84.6% (95% C.I. 69.5 - 94.1%), while specificity was 100% (95% C.I. 97.6 - 100%). The sensitivity of the ELISA (94.9%, 95% C.I. 82.7 – 99.3%) was comparable with previous evaluations. In experimentally infected dogs, earliest positive results with the Angio Detect™ Test were observed 9 weeks post inoculation and 5 weeks later all sera were Angio Detect™ Test positive. After anthelmintic treatment, seropositive dogs turned negative again within 3 to 7 weeks after treatment. The evaluation of the colour intensity of the test strips confirmed the delay of approximately 3-4 weeks for antigen detection by the Angio Detect™ Test compared to the ELISA and its correlation with the time after infection.

Conclusions: This study provided evidence of a good sensitivity and a very high specificity of the rapid device Angio Detect™ Test for detection of circulating A. vasorum antigen in dogs with suspected canine angiostrongylosis, representing a very simple and useful tool to be broadly applied in veterinary practices. The rapid detection of infected dogs is a key point for initiating an indispensable and urgent therapy.

Keywords: Angiostrongylus vasorum, Dogs, Serological diagnosis, Rapid device test, Antigen detection, Specificity, Sensitivity

Background
Canine angiostrongylosis is caused by the metastrongylid Angiostrongylus vasorum, manifesting mainly in respiratory signs, coagulopathies and neurological signs [1], and can be fatal if left untreated [2-4]. Due to this high variability and the often chronic and subtle course of the infections, the diagnosis of this disease is particularly challenging. However, early and correct diagnosis are of paramount importance to prevent the onset of severe pathological changes, considering that they can be present even in the absence of clinical signs [5]. Adult stages of A. vasorum are localized in the pulmonary artery and the right heart of dogs, foxes and some other carnivores, while snails and slugs act as intermediate hosts [6]. The currently most employed method for diagnosis of A. vasorum in dogs is the isolation of first stage larvae excreted with faeces applying larval migration techniques such as the Baermann-Wetzel method, followed by microscopic identification of the larvae [7]. However, during prepatency or
in case of low worm burdens with potential intermittent larval excretion [5,8] or when analysing posted faecal samples that arrive with delay at the laboratory and therefore contain inactive or dead larvae, the test may be negative. Also, differentiation from other lung worm larvae such as *Crenosoma vulpis* and *Filaroides* spp. or larvae that may be present in delayed samples (i.e. of *Strongyloides* and/or hookworms) or in samples collected from the ground (containing free-living or plant parasitic nematodes) needs experienced staff, in order to avoid misdiagnosing. This latter consideration is valid also for FLOTAC, a technique based on the counting of larvae in chambers after spinning faecal samples onto a surface and suggested to improve diagnosis of *A. vasorum* especially when larval migration techniques cannot be used [9].

Attempts for the development of serological methods for detection of *A. vasorum* – infections in dogs started already in 1971 with the detection of specific antibodies against *A. vasorum* [10], and were succeeded by follow-up studies in experimentally infected dogs [11,12]. The identification of immunogenic antigens of different molecular weight in crude extracts of adult female worms in the humoral response of infected dogs [13] was followed by the identification of stage-specific antigens also based on sera of dogs experimentally infected with *A. vasorum* [14]. In a consequent approach, Verzberger-Epshtein et al. [15] used polyclonal rabbit antiserum directed against whole adult worms for the detection of circulating *A. vasorum* antigen. Cross-reactions against several helminths and in particular against *C. vulpis*, a lung worm known to be widely distributed, were evaluated in a sandwich-ELISA, showing high specificity and sensitivity. However, samples from animals infected with *Dirofilaria immitis*, another nematode residing in the heart of definitive hosts in the adult stage and known to produce circulating antigens [16] were not investigated. These both potentially fatal canine nematodes are present in overlapping endemic areas in Europe, and this has to be accounted for when applying diagnostic methods in suspect cases or when analysing dogs in non-endemic areas with travel history. In fact, cross-reactions of sera from dogs infected with *A. vasorum* were detected using commercially available *D. immitis* test kits [17], confirming that antigens of *A. vasorum* and *D. immitis* may share epitopes responsible for potential cross-reactions in antigen detection tests. This fact has been considered during the development of another ELISA developed for the detection of circulating antigen of *A. vasorum* applying a monoclonal antibody which did not react with *D. immitis* excretory/secretory (E/S) antigen [18]. The use of this species-specific monoclonal antibody with polyclonal rabbit antibodies directed against *A. vasorum* adult E/S antigen allowed the detection of circulating antigens with a high specificity (94.0%) and sensitivity (95.7%). In further sandwich-ELISAs, other monoclonal antibodies were used to select potentially diagnostic antigens isolated from the *A. vasorum* adult somatic antigen for the detection of specific antibodies against *A. vasorum*. Also these ELISAs showed high sensitivity (81%) and specificity (98%), and have been validated [19] and used for seropodiemological studies [20-22].

Based on these developments, a rapid device for the serological detection of circulating antigen has been designed (Angio Detect™ Test, IDEXX Laboratories, Westbrook, Maine, USA). The test is intended for routine in-clinic diagnosis, since the rapid detection of infected dogs is a key point for initiating an indispensable and urgent therapy. The aim of this study was to evaluate this newly developed in-clinic test.

**Methods**

Sera from naturally and experimentally infected dogs have been used for determination of sensitivity and specificity. In detail:

a) 39 serum samples from naturally infected dogs were used for determination of sensitivity: 28 sera from Switzerland from dogs suspected for canine angiostrongylosis and confirmed positive by microscopic identification of first stage larvae of *A. vasorum* after Baermann-Wetzel technique [7] at the Institute for Parasitology in Zurich, and 13 sera from Germany from previous lungworm studies [18,23], were also confirmed positive by larval analysis.

b) Sera from 28 dogs experimentally inoculated with a European isolate [5,18,24] and with known worm burden were used for a temporal follow up starting from samples before inoculation and samples collected 21, 35, 49, 62 (± 1, n = 28), 76 (± 1, n = 20), 91 (+1, n = 15) and 97 (n = 4) days post inoculation (dpi). Two dogs were followed-up until 286 dpi. Baermann-Wetzel results of these dogs were available (all dogs became patent within 47-55 dpi).

c) The sera of a further 10 experimentally inoculated dogs were followed up from before inoculation and before and after anthelmintic treatment. Treatment was performed between 81 and 110 dpi and consisted of the administration of imidaclorip 10 mg/kg body weight (BW)/moxidectin 2.5 mg/kg BW spot-on. This dose corresponds to 0.1 ml product (Advocate®) per kg BW. Larval counts were also available for these dogs (all dogs became patent within 47-53 dpi).

d) Cross-reactivity was tested with a total of 121 samples from dogs with proven parasitic infections. In detail, sera of dogs experimentally infected with *Toxocara canis* (n = 21) [25] or *Ancylostoma caninum* (n = 4, from the Institute of Parasitology, University of Veterinary Medicine, Hannover), of German [23] and Swiss dogs naturally infected with
Crenosoma vulpis (n = 18) and (Czech) dogs positive for Oslerus osleri (n = 3) diagnosed by the presence of L1 in faeces, of Italian dogs positive for Eucoleus aerophilus (syn. Capillaria aerophila) eggs (n = 6) detected by coproscopy after flotation and confirmed by PCR-coupled sequencing [26], of dogs positive for Dirofilaria immitis (n = 28) diagnosed by the presence of circulating antigen (DiroCHEK®, Symbiotics, San Diego, USA) and/or microfilariae (which were characterized with the acid phosphatase stain), of dogs positive for Dirofilaria repens (n = 20), Acantocheilonema reconditum (n = 10) or Dipetalonema dracunculoides (n = 10), diagnosed by the presence of microfilariae which were characterized with the acid phosphatase stain and/or length measurements and/or PCR [27], were tested. Also one dog with contemporaneous presence of A. reconditum and D. repens, and two dogs diagnosed with a double infection with A. vasorum and C. vulpis were tested.

e) Finally, 10 control sera of selected dogs negative by Baermann-Wetzel analysis were tested.

All sera were tested with the Angio Detect™ Test (IDEXX Laboratories), a lateral flow immunochromatography test which includes a positive control field. Tests were performed by experienced laboratory staff following the manufacturer’s direction and within the indicated expiry dates. Results were semiquantitatively evaluated based on the colour intensity (+ = slight but visible coloration, ++ = good visible coloration, +++ = intensive coloration, see Figure 1). Furthermore, sera were also tested with the ELISA for detection of circulating antigen of A. vasorum [18]. For a semiquantitative comparison with the results of the Angio Detect™ Test, optical density (OD) values were graduated into negative (< 0.159), + (0.159 - 0.350), ++ (0.351 - 0.800) and +++ (> 0.800) results.

Excel 2007 for Windows (Microsoft Corporation, Redmond, USA) was used to calculate the prevalence values, means and standard deviations (SD). Sensitivities were calculated dividing the number of seropositive animals by the total number of infected animals, while specificities were calculated dividing the number of seronegative animals by the total number of uninfected animals tested. Exact binomial 95% confidence intervals (95% C.I.) for means of binomial variables were calculated with unweighted data according to the method of Clopper and Pearson [28].

All institutional and national guidelines for the care and use of laboratory animals were followed. Experiments with dogs were carried out with facility-born animals at the experimental units of the Vetsuisse Faculty in Zurich upon approval by the Cantonal Veterinary Office of Zurich (permission numbers 25/2006, 26/2007, 10/2008, 13/2008, 185/2008).

Results

Serological Angio Detect™ Test results obtained from dogs naturally infected with A. vasorum are shown in Table 1. A total of 33 out of 39 Baermann-positive dogs were positive, resulting in a sensitivity of 84.6% (95% C.I. 69.5 - 94.1%). In comparison, 37 out of 39 of the sera were also confirmed seropositive for detection of circulating antigen by ELISA (sensitivity 94.9%, 95% C.I. 82.7 – 99.3%). Two out of 6 sera which were negative with the Angio Detect™ Test were also ELISA negative, while the other 4 sera were in the lower OD category (+, OD = 0.159-0.350) of positive results, above the cut-off value.

The serological follow-up of dogs experimentally inoculated with A. vasorum showed that seropositivities detected by Angio Detect™ Test followed seropositivities detected by ELISA in time (Table 2). The earliest positive results with the Angio Detect™ Test were observed at 9 weeks post inoculation: 20 out of 28 sera were ELISA-positive, and five of them were also Angio Detect™ Test
positive. At 11 weeks post inoculation, all tested 20 dogs were seropositive with the ELISA, and 10 of them were also Angio Detect™ Test positive. Thirteen weeks post inoculation (91 dpi), 15 sera were tested, and again all were ELISA-positive, and only 4 sera were Angio Detect™ Test negative: of these, 3 were tested again 97 dpi, and they all became Angio Detect™ Test positive. Dogs followed-up for up to 286 dpi remained constantly seropositive. Concerning the intensity of coloration on the test strips, intensity was by trend higher the longer the dogs were infected. In opposition, colour intensity was not correlated with the number of adult worms determined at necropsy (results not shown).

Out of 10 dogs treated at 81-110 dpi with the spot-on solution containing moxidectin/imidacloprid (Advocate®), seven became positive in the Angio Detect™ Test at 49 (1 dog), 62 (1 dog), 76 (2 dogs), 91 (2 dogs) or 104 (1 dog) dpi. They all turned negative again within 3 (3 dogs) to 7 weeks (3 dogs) after treatment, with one exception: one dog in which one worm was still detected at necropsy remained positive in the Angio Detect™ Test (see also Figure 1) and in the ELISA as well until necropsy (6 weeks after treatment), but was negative for larval excretion. Angio Detect™ Test results were confirmed by negative ELISAs within 3-9 weeks after treatment (after 3, 5, 7 and 9 weeks in 2, 3, 1, and 2 dog, respectively) and absence of larval excretion within 9-20 days after treatment.

As expected, the sera of the two dogs with simultaneous infection with C. vulpis and A. vasorum were positive. All 121 sera tested for potential cross-reactions were negative with the Angio Detect™ Test, indicating a specificity of 100% (95% C.I. 97.6 - 100%). Also the 10 selected sera with negative Baermann-Wetzel analysis were negative with the Angio Detect™ Test, indicating a specificity of 100% (95% C.I. 74.1-100%).

**Discussion**

The diagnosis of canine angiostrongylosis is particularly challenging, since clinical signs may be highly variable. Based on studies with experimentally infected dogs, respiratory signs can be detected approximately seven weeks post inoculation [24], more or less simultaneously with the onset of patency and detection of circulating antigen by ELISA [18]. However, very frequently early clinical signs may not be visible and the onset of potentially fatal coagulopathies may occur [29-32], which may be associated with chronic infections [24,30,33]. Since the first description of A. vasorum in France in the 19th century [34], endemic foci have been described in different European countries, and in the last decades an increasing number of cases have been reported from Europe [21] and from North America in the Atlantic Canadian province of Newfoundland and Labrador [35]. A. vasorum was detected in dogs and in foxes, with potential for expansion and establishment in other areas [36]. This increase of reported cases may be attributed to the increased detection methods, but also to an increase in the distribution range of the definitive host, the red fox (Vulpes vulpes), and to an increasing number of dogs infected with the parasite [37].

Table 1 Serological results of 39 dogs naturally infected with *Angiostrongylus vasorum* (as diagnosed with positive Baermann-Wetzel analysis) tested with the Angio Detect™ Test (IDEXX Laboratories) and with an ELISA for detection of circulating antigen [18]

| Antigen-ELISA² | Angio- detect™ | Negative | + | ++ | +++ | Total |
|----------------|----------------|----------|---|----|-----|-------|
| (< 0.159)      | (0.159-0.350)  | (0.351-0.800) | >0.800 |     |     |       |
| Negative       | 2              | 4        | 6 |    |     | 12    |
| +              | 1              | 3        | 4 |    |     | 8     |
| ++             | 8              | 8        |    |     |     | 16    |
| +++            | 21             | 21       |    |     |     | 42    |
| Total          | 2              | 4        | 1 | 32  | 39  |       |

¹negative = no coloration, + = slight, visible coloration, ++ = good visible coloration, +++ = intensive coloration (see Figure 1).
²ELISA-results categorized based on optical density values (given in brackets).

Table 2 Serological results of dogs experimentally inoculated with *Angiostrongylus vasorum* tested with the Angio Detect™ Test (IDEXX Laboratories) and with the ELISA for detection of circulating antigen [18]

| Weeks after inoculation | No. of dogs | Antigen-ELISA-positive | Angio Detect™ Test -positive |
|------------------------|-------------|------------------------|-----------------------------|
|                         | n           | n                      | % (95% CI)                  | n                         | % (95% CI)                  |
| 0                      | 28          | 0                      | 0 (0-10.1)                  | 0                         | 0 (0-10.1)                  |
| 3                      | 28/9¹       | 0                      | 0 (0-10.1)                  | 0                         | 0 (0-28.3)                  |
| 5                      | 28/9¹       | 6                      | 21.4 (8.3-41.0)             | 0                         | 0 (0-28.3)                  |
| 7                      | 28          | 11                     | 39.3 (21.5-59.4)            | 0                         | 0 (0-10.1)                  |
| 9                      | 28          | 20                     | 71.4 (51.3-86.8)            | 5                         | 17.9 (61.3-36.9)            |
| 11                     | 20          | 20                     | 100 (86.1-100)              | 10                        | 50.0 (27.2-72.8)            |
| 13                     | 15          | 15                     | 100 (81.9-100)              | 11                        | 73.3 (44.9-92.2)            |
| 14                     | 8           | 8                      | 100 (68.8-100)              | 8                         | 100 (68.8-100)              |
| 15-41                  | 2           | 2                      | 100 (22.4-100)              | 2                         | 100 (22.4-100)              |

¹only 9 out of the 28 dogs were tested with the Angio Detect™ Test.

Baermann-Wetzel confirmed patency in all dogs within 47-55 dpi.
to increased disease awareness from the veterinary and from the pet owner’s side as well, but also due to available new diagnostic techniques such as molecular tools [37-39] or ELISAs, contributing to increased knowledge and diagnosis of *A. vasorum*. Molecular techniques are particularly recommended for prevalence studies in intermediate mollusc hosts [40] and for confirmative diagnosis in dogs or foxes [37], while ELISAs have been used for mass-screening in large epidemiological investigations [21,22]. Additionally, as previously shown, the specific detection of circulating *A. vasorum* antigen by ELISA also represents a valid alternative for reliable diagnosis and for follow-up investigations after anthelmintic treatment [18].

Both tests, Angio Detect™ Test and the ELISA for detection of circulating antigen, are intended to diagnose an active *A. vasorum* infection, and sensitivity and specificity on naturally infected dogs of this study were 84.6% (95% C.I. 69.5 - 94.1%) and 94.9% (95% C.I. 82.7 – 99.3%), for the rapid device and the ELISA, respectively. The sensitivity of the ELISA determined in this study was absolutely comparable with the results previously obtained with sera of 23 naturally infected dogs (95.7%, 95% CI 78.1-99.9% [18]).

Other commercially available tests based on the same principle as the Angio Detect™ Test (lateral flow immunochromatography), in particular tests for the detection of circulating antigen of *D. immitis*, were frequently showing lower sensitivities compared to plate ELISA assays which involve multiple steps including signal amplification. In a study performed with sera of dogs harbouring low heartworm burdens, sensitivities of the 3 tested lateral flow immunochromatography tests varied between 52 and 71%, while three test kits based on ELISA had sensitivities between 67-71% [41]. In another study performed with dog sera collected in Argentina, two kits based on lateral flow immunochromatography and one on membrane ELISA all had a sensitivity of 76% [42]. Lower sensitivities of test kits based on lateral flow immunochromatography compared with ELISAs has also been reported for other parasites such as *Leishmania* infections in dogs, with sensitivities varying between and 53% and 64% (asymptomatic dogs), and 76% and 97% (symptomatic dogs) for test kits and ELISAs, respectively [43]. In this context, the calculated sensitivity of 84.6% of the Angio Detect™ Test can be considered as high and is additionally confirmed by detection of an *A. vasorum* infection in a treated dog which was negative for larval excretion but in which one worm was still detected at necropsy. In fact, as already mentioned, the Baermann-Wetzel technique can be false negative also after prepatency.

Four out of 6 sera of naturally infected dogs which were negative with the Angio Detect™ Test were allocated to the “low positive” category in the ELISA; possibly, these dogs would have become seropositive with the Angio Detect™ later. Differences of sensitivities between Angio Detect™ Test and ELISA for detection of *A. vasorum* infections are illustrated more in detail by the results obtained when testing experimentally infected dogs. With increasing time after inoculation, seropositivity increased for both tests: detection of circulating antigen with the ELISA started from 5 weeks after inoculation and all tested dogs were positive starting from 11 weeks after inoculation. In contrast, first seropositive results with the Angio Detect™ Test were obtained starting from 9 weeks after inoculation and were positive for all dog sera at 14 weeks after inoculation. The evaluation of the colour intensity of the test strips confirmed the delay of approximately 3-4 weeks for antigen detection by the Angio Detect™ Test and its correlation with the time after infection, independent of the worm burden detected at necropsy. However, this delay could be of minor importance for prevention of potentially deadly cases: as previously shown, despite heavy *A. vasorum* infection load and pulmonary changes, only mild haematological changes were observed in experimentally infected dogs up to 13 weeks after inoculation [24], indicating that fatal coagulopathies may occur only afterwards. At this time, sensitivity of the Angio Detect™ Test increased up to 100%, and was additionally complemented with a very high specificity of 100% (95% C.I. 97.6-100%).

Interestingly, both ELISA and Angio Detect™ Test were negative for two dogs diagnosed positive by Baermann-Wetzel technique. A tentative explanation of this result could be the formation of antigen-antibody complexes, which may inhibit detection of antigen in some canine samples, as it has been shown with *D. immitis* [44,45]. Similarly, during the evaluation of the antigen-ELISA [18], the follow-up of an experimentally infected dog harbouring 165 adult *A. vasorum* at necropsy evidenced OD values which were constantly under the cut-off value, except at 35 dpi, possibly due to relevant amounts of circulating antigens and antibodies forming antigen-antibody complexes. In order to test this hypothesis, we re-tested the serum of this latter dog and some of the sera of the study presented here, for which an adequate amount of serum was available after pretreatment with heat, as previously described [46]. One out of two sera of the naturally infected dogs negative for both, the antigen-ELISA and the Angio Detect™ Test (see Table 1), turned out positive in both tests. A further 12 sera of naturally and experimentally infected dogs with contradictory results (antigen-ELISA positive and Angio Detect™ Test negative) were retested after heat treatment: 9 of them were then also positive in the Angio Detect™ Test (data not shown). This indicates that pretreating the sera may increase the sensitivity of the test. However, there were still sera of dogs with proven
experimental infections that remained negative, in opposition to the ELISA result.

Currently, canine angiostrongylosis is diagnosed with analysis of faeces by the Baermann-Wetzel method, having the disadvantage that larval migration is usually performed overnight and additionally samples are often sent to a laboratory, causing a delay in diagnosis. Considering that the Angio Detect™ Test can be directly used in veterinary practices for suspect dogs, this test represents a valid alternative for immediate diagnosis and subsequent concerted treatment of angiostrongylosis. Certainly, coproscopic methods still remain an option in suspect dogs and with negative Angio Detect™ Test results, and represent the only option for diagnosis of other lung worms such as *Crenosoma vulpis* or *Eucoleus aerophilus*.

**Conclusions**

This study provided evidence of a good sensitivity and a very high specificity of the rapid device Angio Detect™ Test for detection of circulating *A. vasorum* antigen in dogs with suspected canine angiostrongylosis. As the test is conceived for in clinic-use, its use is simple and easy and can be promptly and broadly applied, representing a very useful tool to be used in veterinary practices.

**Competing interests**

MS and PD were involved in the design of the Angio Detect™ Test.

**Authors’ contributions**

MS participated in the design of the study, collected the samples, carried out the diagnostic assays and drafted the manuscript. KS carried out the diagnostic assays. TJN collected the samples and contributed to the draft of the manuscript. SL collected the samples and carried out the diagnostic assays. PD participated in the design of the study and the draft of the manuscript. All authors have read and approved the final manuscript.

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**References**

1. Koch J, Willeisen JL. Canine pulmonary angiostrongylosis: an update. *Vet J* 2009, 179(3):348–359.
2. Denk D, Matisek N, Just FT, Herrmanns W, Baiker K, Herbach N, Steinberg T, Fischer A: Disseminated angiostrongylosis with fatal cerebral haemorrhages in two dogs in Germany: a clinical case study. *Vet Parasitol* 2009, 160(1–2):100–108.
3. Staetble S, Ochs H, Steffen F, Naegeli F, Borel N, Sieber-Ruckstuhl N, Deplazes P: Autochthonous infections with *Angiostrongylus vasorum* in dogs in Switzerland and Germany (in German). Schweiz Arch Tierheilkd 2005, 147(3):121–127.
4. Yomawaki Y, McGarry JV, Denk D, Dukes-McEwan J, Macdonald N, Mas A, McConnell F, Tartton B, Valentine GC, Wayne J, et al: Emerging canine angiostrongylosis in northern England: five fatal cases. *Vet Rec* 2009, 164(5):149–152.
5. Schnyder M, Fahrion A, Osent P, Kohler L, Webster P, Heine J, Deplazes P: Larvicidal effect of imidacloprid/moxidectin spot-on solution in dogs experimentally inoculated with *Angiostrongylus vasorum*. *Vet Parasitol* 2009, 166:326–332.
6. Gilhoun J: Recherches sur le cycle évolutif du Strondge des vaisseaux du chien. *Bull Acad Vét* 1963, 64:431–442.
7. Lehrbuch der Parastologie für die Tiermedizin. 3rd edition Edited by Deplazes P, Eckert J, Samson-Himmelstjerna V, Zahner H. Stuttgart: Enke Publishers; 2013.
8. Oliveira-Junior SD, Barcante JM, Barcante TA, Dias SR, Lima WS: Larval output of infected and re-infected dogs with *Angiostrongylus vasorum* (Baillet, 1866) Kamensky, 1905. *Vet Parasitol* 2006, 141(1–2):101–106.
9. Schnyder M, Maurelli MP, Morgogline ME, Kohler L, Deplazes P, Torgerson P, Cirigoli G, Rinaldi L: Comparison of faecal techniques including FLOTAC for copromicroscopic detection of first stage larvae of *Angiostrongylus vasorum*. *Parasitol Res* 2011, 109:63–69.
10. Gilhoun J, Benex J, Mishra GS: First attempts of immunological diagnosis of canine angiostrongylosis due to *Angiostrongylus vasorum* (in French). *Vet Parasitol* 2011, 181:220–228.
11. Costa JO, Lima DOS Santos W, Nascimento E: Imunodiagnostico por ELISA da angiostrongilose canina experimental. *Aby Bras Med Zootec* 1996, 48(4):491–495.
12. Cury MC, Lima WS, Vitor RW: Enzyme-linked immunosorbent assay (ELISA) for the diagnosis of *Angiostrongylus vasorum* (Baillet, 1866) infection in dogs. *Rev Med Vet* 1996, 147(7):525–530.
13. Cury MC, Guimarães MP, Lima WS, Vitor RW: Western blot analysis of the humoral response of dogs experimentally infected with *Angiostrongylus vasorum* (Baillet, 1866). *Vet Parasitol* 2002, 106(1–3):83–87.
14. De Oliveira Vasconcelos V, Wagner De Almeida VR, Dos Santos Lima W: Identification of stage-specific proteins of *Angiostrongylus vasorum* (Baillet, 1866) Kamensky. *Vet Parasitol* 2008, 152(1–2):389–395.
15. Verzicer-epstein I, Markham RJ, Sheppard JA, Stryin H, Whitney H, Conboy GA: Serologic detection of *Angiostrongylus vasorum* infection in dogs. *Vet Parasitol* 2008, 151(1):53–60.
16. Ehrenberg JP, Tamashiro WK, Scott AL: *Dirofilaria immitis*: identification and characterization of circulating parasite antigens. *Exp Parasitol* 1987, 63(2):205–214.
17. Schnyder M, Deplazes P: Cross-reactions of sera from dogs infected with *Angiostrongylus vasorum* in commercially available *Dirofilaria immitis* test kits. *Parasit Vectors* 2012, 5:258.
18. Schnyder M, Tanner I, Webster P, Barutzki D, Deplazes P: An ELISA for sensitive and specific detection of circulating antigen of *Angiostrongylus vasorum* in serum samples of naturally and experimentally infected dogs. *Vet Parasitol* 2011, 179:152–158.
19. Schucan A, Schnyder M, Tanner I, Barutzki D, Traversa D, Deplazes P: Detection of specific antibodies in dogs infected with *Angiostrongylus vasorum*. *Vet Parasitol* 2012, 185:216–224.
20. Guardone L, Schnyder M, Macchioni F, Deplazes P, Magi M: Serological detection of circulating *Angiostrongylus vasorum* antigen and specific antibodies in dogs from central and northern Italy. *Vet Parasitol* 2013, 192(1–3):192–198.
21. Schnyder M, Schaper R, Bilbrough G, Morgan ER, Deplazes P: Seropidemiological survey for canine angiostrongylosis in dogs from Germany and the UK using combined detection of *Angiostrongylus vasorum* antigen and specific antibodies. *Parasitology* 2013, 140(1):142–1450.
22. Schnyder M, Schaper R, Pantchev N, Kowalska D, Szvedko A, Deplazes P: Serological detection of circulating *Angiostrongylus vasorum* antigen and parasite-specific antibodies in dogs from Poland. *Parasitol Res* 2013, 112(Suppl 1):109–117.
23. Barutzki D, Schaper R: Natural infections of *Angiostrongylus vasorum* and *Crenosoma vulpis* in dogs in Germany (2007–2009). *Parasitol Res* 2009, 105(Suppl 1):S39–S48.
24. Schnyder M, Fahinion A, Rond A, Oriss P, Webster P, Kranjc A, Glaus T, Deplazes P. Clinical laboratory and pathological findings in dogs experimentally infected with Angiostrongylus vasorum. Parasitol Res 2010, 107:1471–1480.

25. Fahinion AS, Staebler S, Deplazes P. Patent Toxocara canis infections in previously exposed and in helminth-free dogs after infection with low numbers of embryonated eggs. Vet Parasitol 2008, 152:1–2:108–115.

26. Traversa D, Di Cesare A, Lia RP, Castagna G, Meloni S, Heine J, Strube K, Millili P, Oronero D, Neckes O, et al. New insights into morphological and biological features of Capillaria aerophila (Trichocephalida, Trichuridae). Parasitol Res 2011, 109(Supp 1):597–5704.

27. Magnis J, Lorentz S, Guardone L, Grimm F, Magi M, Naucke TJ, Deplazes P. Morphometric analyses of canine blood microfilariae isolated by the Knott’s test enables Dirofilaria immitis and D. repens species-specific and Acanthocheilonema (syn. Dipetalonema) genus-specific diagnosis. Parasit Vectors 2013, 6:48.

28. Clopper CJ, Pearson ES. The use of confidence or fiducial limits illustrated in the case of the binomial. Biometrika 1934, 26:404–413.

29. Cury MC, Lima WS, Guimaraes MP, Cavalcio MG. Hematological and coagulation profiles in dogs experimentally infected with Angiostrongylus vasorum (Baillet, 1866). Vet Parasitol 2002, 104(2):139–149.

30. Ramsey JK, Littlewood JD, Dunn JK, Heritage ME. Role of chronic disseminated intravascular coagulation in a case of canine angiostrongylosis. J Vet Rec 1996, 138(15):360–363.

31. Schelting CG, Greene CE, Prestwood AK, Tsang VC. Coagulation abnormalities associated with acute Angiostrongylus vasorum infection in dogs. Am J Vet Res 1986, 47(12):2669–2673.

32. Weissmann A, Lu D, Lamb CR, Smyth B, Mantis P, Chandler K, Boag A, Cherubini GB, Cappello R. Brain and spinal cord haemorrhages associated with Angiostrongylus vasorum infection in four dogs. Vet Rec 2006, 158(25):858–863.

33. Schmitz S, Moritz A. Chronic disseminated intravascular coagulopathy in a dog with lung worm infection (in German). Schweiz Arch Tierheilkd 2009, 151(6):281–286.

34. Serres E. Entozoaires trouvés dans l’oreille droite, le ventricule correspondant et l’artère pulmonaire d’un chien. J Vet Med 1854, 7:70.

35. Conboy GA. Canine angiostrongylosis: the French heartworm; an emerging threat in North America. Vet Parasitol 2011, 176(4):382–389.

36. Morgan ER, Jefferies R, Krajewski M, Ward P, Shaw SE. Canine pulmonary angiostrongylosis: the influence of climate on parasite distribution. Parasitol Int 2009, 58(4):406–410.

37. Al-Sabi MN, Deplazes P, Webster P, Willesen JL, Davidson RK, Kapel CM. PCR detection of Angiostrongylus vasorum in faecal samples of dogs and foxes. Parasitol Res 2010, 107:135–140.

38. Jefferies R, Morgan ER, Shaw SE. A SYBR green real-time PCR assay for the detection of the nematode Angiostrongylus vasorum in definitive and intermediate hosts. Vet Parasitol 2009, 166(1–2):112–118.

39. Traversa D, Guglielmini C. Feline aelurostrongylosis and canine angiostrongylosis: a challenging diagnosis for two emerging verminous pneumonia infections. Vet Parasitol 2008, 157(3–4):163–174.

40. Ferndreuth T, Kapel CM, Webster P, Al-Sabi MN, Gronvold J. The occurrence of Angiostrongylus vasorum in terrestrial slugs from forests and parks in the Copenhagen area, Denmark. J Helminthol 2009, 83:279–285.

41. Courteney CH, Zeng Q. Comparison of heartworm antigen test kit performance in dogs having low heartworm burdens. Vet Parasitol 2001, 96(4):317–322.

42. Vezzani D, Fontanarossa VF, Eiras DF. Are antigen test kits efficient for detecting heartworm-infected dogs at the southern distribution limit of the parasite in South America? Preliminary results. Res Vet Sci 2008, 85(1):113–115.

43. Mettler M, Grimm F, Capelli G, Camp H, Deplazes P. Evaluation of enzyme-linked immunosorbent assays, an immunofluorescent-antibody test, and two rapid tests (immunochromatographic-dipstick and gel tests) for serological diagnosis of symptomatic and asymptomatic Leishmania infections in dogs. J Clin Microbiol 2005, 43(11):5515–5519.

44. Brunner CJ, Hendrix CM, Blaibourn BL, Hanrahan LA. Comparison of serologic tests for detection of antigen in canine heartworm infections. J Am Vet Med Ass 1988, 192(10):1423–1427.

45. Matsumura K, Kazuta Y, Endo R, Tanaka K, Inoue T. Detection of circulating immune complexes in the sera of dogs infected with Dirofilaria immitis, by Clq-binding enzyme-linked immunosorbent assay. J Helminthol 1986, 60(3):239–243.

46. Weil GJ, Malone MS, Powers KG, Blair LS. Monoclonal antibodies to parasite antigens found in the serum of Dirofilaria immitis-infected dogs. J Immunol 1985, 134(2):1185–1191.