Angiostrongylus vasorum in the eye: new case reports and a review of the literature

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Abstract

Background: Nematodes of the genus Angiostrongylus are important causes of potentially life-threatening diseases in several animal species and humans. Angiostrongylus vasorum affects the right ventricle of the heart and the pulmonary arteries in dogs, red foxes and other carnivores. The diagnosis of canine angiostrongylosis may be challenging due to the wide spectrum of clinical signs. Ocular manifestations have been seldom reported but have serious implications for patients.

Methods: The clinical history of three cases of infection with A. vasorum in dogs diagnosed in UK, France and Italy, was obtained from clinical records provided by the veterinary surgeons along with information on the diagnostic procedures and treatment. Nematodes collected from the eyes of infected dogs were morphologically identified to the species level and molecularly analysed by the amplification of the nuclear 18S rRNA gene.

Results: On admission, the dogs were presented with various degrees of ocular discomfort and hyphema because of the presence of a motile object in the eye. The three patients had ocular surgery during which nematodes were removed and subsequently morphologically and molecularly identified as two adult males and one female of A. vasorum.

Conclusions: Three new cases of canine ocular angiostrongylosis are reported along with a review of other published clinical cases to improve the diagnosis and provide clinical recommendation for this parasitic condition. In addition, the significance of migratory patterns of larvae inside the host body is discussed. Veterinary healthcare workers should include canine angiostrongylosis in the differential diagnosis of ocular diseases.

Keywords: Angiostrongylus vasorum, Lungworm, Ocular infection, Eye, Metastrongyloidea, Diagnosis, Snails

Background

Nematodes of the genus Angiostrongylus Kamensky, 1905 (Strongylida, Angiostrongylidae) are important because of their life-threatening potential in several animal species and humans [1]. Helminths within the superfamily Metastrongyloidea are usually known as “lungworms” because of their localisation in the lungs and associated blood vessels in the definitive host [2]. Angiostrongylus spp. develop in and are transmitted by gastropods (i.e. snails and slugs) which act as intermediate hosts [1, 3]. Amongst these parasites, Angiostrongylus cantonensis and Angiostrongylus costaricensis infect rodents and, occasionally, humans causing eosinophilic meningitis [4, 5] and abdominal angiostrongylosis [6], respectively. Dogs have been indicated as definitive hosts of A. costaricensis, suggesting that they may act as a reservoir host for this parasite in the domestic environment [7]. Angiostrongylus vasorum may cause severe clinical disease in dogs, red foxes and other carnivores, characterised by respiratory distress [8]. This parasitic infection has a patchy distribution in many parts of the world, including tropical, subtropical and temperate regions (i.e. Europe, Africa, North and South America) [9], and it is apparently expanding in new areas and around well-defined endemic foci [10]. The lack of international surveillance and difficulties in diagnosing A. vasorum impedes collection of data on its spread and global distribution [10–12]. Nevertheless, improved knowledge of parasite biology is required before drawing any definitive conclusion about the significance of the
geographical expansion of *A. vasorum* [8, 9]. For instance, snail-to-snail transmission of infective third-stage larvae (L3) of *Aelurostrongylus abstrusus* (Strongylida, Angiostrongylidae) has been hypothesised as a key example for the spreading of nematodes associated with gastropod-borne diseases in endemic areas [13].

As with other metastrongylid nematodes, *A. vasorum* develops in snails and slugs from first-stage larvae (L1) to infective L3, in approximately 16 days [14]. In the canid definitive hosts, L3 undergo two moults in the abdominal lymph nodes and fifth stage larvae (L5) reach the right ventricle and pulmonary arteries, where they develop into dioecious adult nematodes [2, 15]. Gravid females lay eggs in the bloodstream that hatch in the respiratory system and L1 are passed out in the faeces [2, 15]. Canids may release L1 during their whole life, although the frequency of larval shedding varies over the year [8]. Along with this typical route of infection, frogs (*Rana temporaria*) may also act as intermediate and paratenic hosts of *A. vasorum* [16]. In addition, dogs can be experimentally infected with L3 shed in the environment from the snail *Biomphalaria glabrata* [17].

Clinical diagnosis of canine angiostrongylosis is challenging because of the wide spectrum of clinical signs and because subclinical infections also occur, leading to an underestimation of the true prevalence of infection [11]. Indeed, while respiratory signs are considered the main clinical presentation of the infection by *A. vasorum*, coagulative, cardiovascular and neurological disorders are also described [9]. The clinical picture can be further complicated by the fact that this condition may remain undiagnosed for months or even years [9]. Whilst respiratory and cardiac clinical signs are most commonly associated with *A. vasorum* infection, ocular manifestations have been seldom reported [18]. A better awareness of ocular angiostrongylosis will improve its diagnosis and treatment.

Here we report three new cases of canine ocular angiostrongylosis together with a review of previously published clinical cases.

**Methods**

**Case presentation**

Information on the clinical history of two cases of *A. vasorum* infection in dogs diagnosed in the UK and France (Cases 1 and 2, respectively) was obtained from clinical records provided by the veterinary practitioners along with information on the diagnostic procedures, treatment and outcome. A third case from Italy (Case 3) was also included [19]. In addition, we review cases reported in the international literature between 1913 and 2015, searching in Google Scholar and in the PubMed database the keywords “ocular angiostrongylosis”, “Angiostrongylus vasorum eye” and “Angiostrongylus vasorum ocular”.

**Morphological and molecular identification**

Nematodes collected from the eyes of infected dogs were morphologically identified to the species level based on previous descriptions [20, 21]. In addition, identity of specimens extracted from Case 1 and 2 were confirmed by PCR. Briefly, genomic DNA from adult worms and L1 was extracted using a commercial kit (DNeasy Blood & Tissue Kit, Qiagen, GmbH, Hilden, Germany), in accordance with the manufacturers’ instructions. A nuclear 18S rRNA gene (~1700 bp) was amplified using the following primers (NC18SF1: 5’-AAA GAT TAA GC C ATG CA-3’ and NC5BR: 5’- GCA GGT TCA CCT ACA GAT-3’). The amplicons were purified and sequenced using the Taq Dye Doxy Terminato Cycle Sequencing Kit (v.2, Applied Biosystems, Foster City, CA) in an automated sequencer (ABI-PRISM 377). Sequences were compared with those available in the GenBank database by Basic Local Alignment Search Tool (BLASTn http://blast.ncbi.nlm.nih.gov/Blast.cgi).

**Ethics statement**

All medical procedures were carried with the owner’s approval. Nematodes were collected by veterinarians working in private clinics and sent to the Laboratory of Parasitology (University of Bari, Italy) for diagnostic purposes.

**Results**

**Case 1**

A 21-month-old female pug was referred to a veterinary ophthalmologist with a one-day history of trauma to the right eye. The dog lived in a rural environment, on the edge of woods in the Greater Manchester area, in the vicinity of many animals (i.e. horses, sheep, pigs, chickens, and other dogs) and of a river, where the dog is usually walked. At the clinical visit, panuveitis with hyphema and fibrin deposition in the right eye were observed; partial examination of the posterior segment showed no evidence of retinal detachment or posterior haemorrhages. Rebound tonometry revealed acute ocular hypertension with intraocular pressure of 30 mmHg in the right eye (20 mmHg in the left eye); however, a dazzle reflex was still present. No intraocular parasite was detected during the initial examination and there was no evidence of a systemic disease. The following day, the dog was sedated and a free floating worm was found in the anterior chamber of the right eye. The worm was aspirated via anterior chamber paracentesis using a 21 gauge needle and attached 2 ml syringe, placed in ethanol and subsequently morphologically and molecularly identified. An intracameral injection of 25 μg of tissue plasminogen activator and 0.1 mg of adrenaline was injected into the right anterior chamber to facilitate fibrinolysis, induce mydriasis and minimise further haemorrhage. Medical treatment consisted of chloramphenicol
drops qid, brinzolamide drops qid, prednisolone acetate drops qid, nepafenac drops qid, cephalexin 15 mg/kg bid and prednisolone 0.5 mg/kg SID per os. Milbemycin oxime 12.5 mg combined with praziquantel 125 mg (Milbemax®, Novartis Animal Health) was prescribed once a week for 4 weeks, then once monthly long term.

The parasite was broken at the level of the anterior extremity measuring the two main pieces 2.13 mm and 4.51 mm in length; the width was 0.118, 0.28 and 0.182 mm at the anterior, middle and posterior ends, respectively. The nematode had a moderately dilated cuticle at the cephalic extremity, with a small buccal aperture (Fig. 1a). The oesophagus measured 0.219 mm in length, and the excretory pore was located at 0.410 mm from oesophagus. The posterior extremity was ventrally curved, with a small copulatory bursa with well-developed bursal rays (Fig. 1b). Spicules appeared yellowish and strongly sclerotised and transversally striated and measured 0.456 mm and 0.459 mm in length (Fig. 1c, d). A slightly sclerotised ellipsoidal gubernaculum was located close to the terminal end of the spicules. The nematode was identified as a male of *A. vasorum*. Further, the 18S rDNA sequence obtained from the adult nematode displayed 100 % identity to the nucleotide sequence of *A. vasorum* [GenBank: AJ920365].

A follow-up 2 weeks later, revealed complete resolution of the anterior uveitis and hyphema. Two 1 mm white opacities and one small resorbing hemorrhage were detected retrolenticularly in the anterior vitreous. Further examinations, investigations and treatment were declined by the owners. A follow-up by telephone was performed 20 months later and the owner reported no further evidence of ocular or systemic clinical signs.

**Case 2**

A 2-year-old male Cavalier King Charles Spaniel dog was referred to a private veterinary clinic in Paris (France) due to persistent blepharospasm and epiphora. The dog that lived in the city centre was walked in a forest nearby. On clinical ophthalmological examination the dog showed prolapse of the nictitating membrane, photophobia on the left eye, iris hyperaemia and corneal oedema. The intraocular pressure was 14 mmHg in the left eye and 16 mmHg in the right eye, the fluorescein test was negative and the Schirmer test showed a slightly increased lacrimation. A thread-like organism was noticed in the anterior chamber of the left eye (Fig. 2), which was very motile under light stimuli. An additional movie file clearly shows this in more detail (see Additional file 1).

Removal of the parasite was performed by anterior chamber paracentesis and the nematode was morphologically and molecularly processed. The specimen was identified as an unfertilised *A. vasorum* female. Briefly, the female nematode measured 16 mm in length and 0.2 mm in width; genital ducts were coiled around the reddish intestine, which appeared visible throughout the

**Fig. 1** Case 1. Cephalic extremity of *Angiostrongylus vasorum* (a); Short copulatory bursa featured by well-developed bursal rays (b); Strongly sclerotised spicules with a thin membrane (c); Spicules measured 0.456 mm and 0.459 mm in length (d)
cuticle. However, the nematode was damaged and further morphological features were not evaluated, with the exception of the uteri, which lacked first-stage larvae, and the vulva. Therefore, the dog was subjected to coprological examination for the detection of L1 using the Baermann technique. The 18S rDNA sequences obtained from both L1 collected from the faeces and the female nematode displayed 100 % identity to the nucleotide sequence of *A. vasorum* [GenBank: AJ920365]. The dog lacked any signs of respiratory infection, both previously and during the observation period. The animal was treated with fenbendazol 25 mg/Kg *per os* SID for 3 weeks associated with prednisolone 0.2 mg/Kg.

**Case 3**

A 5-month-old male mixed breed dog was referred to a private veterinary clinic in Rome (Italy) for sudden visual loss. At the admission, the clinical ophthalmological examination revealed corneal oedema and episcleral congestion in the right eye. Examination of the anterior chamber showed diffuse hyphema and complete examination of the posterior segment could not be performed. Clinical diagnosis of anterior uveitis in the right eye was made. Ultrasound of the right eye displayed a blood clot in the anterior chamber. CBC, biochemistry and electrolytes were within the limits. Topical dexamethasone 0.2 %, four times a day, was administered for 2 weeks. Fifteen days following the first examination, the hyphema disappeared and the presence of a free-swimming nematode in the anterior chamber was noticed. Subsequently the dog was anesthetised and the eye was clipped and prepared for surgery. Under surgical microscope the cornea was incised, the anterior chamber was flushed with balanced saline solution (BSS) and the nematode was collected and morphologically identified. Briefly, the parasite presented a smooth cuticle and a slender body with a small buccal aperture (Fig. 3a), and measured 10.5 mm in length and 0.29 mm in width at the middle portion, the anterior and posterior extremities were 0.109 and 0.176 mm in width, respectively (Fig. 3b, c). The oesophagus was 0.219 mm long. The posterior end was ventrally curved, with a short copulatory bursa (Fig. 3d). The subequal spicules measured 0.404 mm and 0.388 mm in length. The nematode was morphologically identified as a male of *A. vasorum*. Unfortunately, due to inadequate preservation of the specimen, the extraction and amplification of genomic DNA was unsuccessful. Stool samples were taken to look for L1 with Baermann technique and tested negative. The animal recovered without complications after the surgery. Topical and systemic antibiotics and steroids were administered for the next 3 weeks. The dog was re-examined at 2 and 6 months post-operatively and ocular examinations were within normal limits.

**Discussion**

Three cases of ocular infection with *A. vasorum* in dogs are described along with clinical presentations and diagnostic procedures. This information is vital to help our understanding of the clinico-pathological picture of canine
ocular angiostrongylosis. Several reports of aberrant migration of *A. vasorum* are described in the scientific literature, with adult worms recovered from left ventricle, pericardial sac, urinary bladder and femoral artery [22–24], and L1 from brain, kidney, liver, muscle, stomach, pancreas and spleen of infected dogs [22, 23, 25]. However, to date, ocular migration of *A. vasorum* has rarely been described. Cases of canine ocular angiostrongylosis have been reported in France [26–28], United Kingdom [18, 29], Denmark [30] and Canada [25] (Table 1), although many infections may be unreported or inadequately described. For instance, in a series of six cases of ocular parasitosis described by [18], dogs presented clinical signs strongly suggestive of *A. vasorum* infection but the aetiology of these conditions was not confirmed by any morphological or molecular identification. Our findings suggest that *A. vasorum* infection should be considered if ophthalmological examination reveals a nematode in the eye even in the absence of other typical clinical signs and L1 are not detected in faeces [11, 18, 28]. In Case 2 as well as in the report by Rosenlund et al. [30], despite the absence of any cardiopulmonary sign, L1 were found in the faeces only after the diagnosis of ocular angiostrongylosis. Conversely, in Case 3 and in the report by Payen [28], L1 were not detected in the faeces. Although in the case described by Payen [28] the dog was presented with overt cardiorespiratory clinical signs, the diagnosis of *A. vasorum* infection was based on the identification of the nematode in the anterior chamber of the eye. As shown in Case 3 and by King et al. [29], dogs were presented with varying degrees of ocular discomfort, and only upon re-examination six to 15 days later, was the presence of a free-swimming nematode detected. This emphasises the importance of clinical follow-up in order to eventually diagnose ocular infection with *A. vasorum*.

Other nematodes have been identified in the eyes of dogs, notably fourth-stage larvae of *Dirofilaria immitis* (Spirurida, Onchocercidae) which have been shown to migrate in the anterior chamber and vitreous body of the eye [31, 32]. In addition, adult *Onchocerca lupi* (Spirurida, Onchocercidae) usually localise in the subconjunctival granulomas and/or in the retrobulbar space of the eye of infected dogs [33, 34], and recently intraocular onchocercosis has also been described in a patient suffering from anterior uveitis [35]. The availability of a diagnostic test for the detection of circulating *A. vasorum* antigens in dogs (IDEXX Angio Detect™) [36] has provided a further tool to assist a definitive parasitological diagnosis. Interestingly, all dogs suffering from canine ocular angiostrongylosis were under the 3 years of age (i.e. 5 months to 3 years), and of the few reports now available in literature, three [28, 29] and in Case 2, involved Cavalier King Charles Spaniel dogs. However, additional epidemiological data are needed for a clear assessment of risk factors (e.g. breed and age) related to the occurrence of canine ocular angiostrongylosis.

![Fig. 3 Case 3. Smooth cuticle and slender body of *Angiostrongylus vasorum* (a); Anterior extremity (b); Posterior extremity (c); Posterior extremity at higher magnification showing a short copulatory bursa (d).](image-url)
How *A. vasorum* larvae reach the ocular tissues of dogs is not clear, although migration may take place by penetration of the corneal surface (cranial-hypodermis route), the surface of the brain and the optic foramen (intracranial-optic foramen route) or through a fibrin sac in the anterior chamber of the eye (corneal route), in a similar manner to those reported for *D. immitis* [37].

The hyphema described in Case 1 and 3 has been associated with the infection as a likely consequence of coagulative disorder or a traumatic effect of the adult nematode on the ocular tissues. This could explain the finding of the adult worm in the anterior chamber only at the second examination. Nevertheless, further clinico-parasitological research is necessary to ascertain the routes of migration and the pathogenesis of ocular angiostrongylosis in dogs. For instance, for many zoonotic helminths affecting the eyes, the parasitic localisation in the ocular tissues or the immune reaction elicited by adults or larval stages in the host are of primary importance in the appearance of overt clinical signs [38].

Milbemycin oxime combined with praziquantel once a week for 4 weeks and fenbendazole (25 mg/kg) for 3 weeks were used to treat *A. vasorum* infection in Case 1 and 2, respectively. Several pharmaceutical options are now available and highly efficacious in treating canine angiostrongylosis, including moxidectin/imidacloprid spot on solution (Advocate<sup>®</sup>, Bayer Animal Health) with a single monthly application [39, 40], and milbemycin oxime in combination with praziquantel (Milbemax<sup>®</sup>, Novartis Animal Health) administered weekly for 4 weeks [41].

**Conclusion**

Aberrant migration of nematodes increases the complexity of the clinico-pathological picture of canine angiostrongylosis, thus an enhanced awareness of clinical conditions caused by *A. vasorum* is imperative for the diagnosis and treatment of this infection. Recognition of the importance of alternative migratory routes of *A. vasorum* in dogs will improve our current understanding of the diagnosis and clinical follow-up of this parasitic condition. For example, veterinary healthcare workers should include canine angiostrongylosis in the differential diagnosis of ocular diseases. Finally, in a review of 484 cases of human eosinophilic meningitis caused by *A. cantonensis*, 47 patients (9.7 %) suffered from ocular disease [42]. In addition, at least 35 patients were further diagnosed with human ocular angiostrongylosis [43]. Therefore, a better appreciation of ocular angiostrongylosis in dogs may assist in our understanding of human ocular angiostrongylosis.

**Table 1 Review of cases described in the literature of canine ocular angiostrongylosis, along with data on the location, clinical presentation, diagnosis and anthelmintic treatment**

| Age, sex and breed | Location | Clinical presentation | Diagnosis | Anthelmintic treatment | Reference |
|--------------------|----------|-----------------------|-----------|------------------------|-----------|
| 8-month-old; n/a; Cavalier King; Charles Spaniel | France | Cough, uveitis, and presence of a motile worm in the anterior chamber of the eye | Faecal and broncho-alveolar washing examination negative for *A. vasorum* larvae. One immature *A. vasorum* female extracted from the eye | Fenbendazole-based treatment for 2 weeks | [28] |
| 14-month-old; female; Cocker Spaniel | Denmark | Epiphora, circumcorneal and conjunctival injection of the episcleral blood vessels, blepharospasm, prolapse of the nictitating membrane miotic and swollen iris. Presence of a motile worm in the anterior chamber of the eye | Numerous *A. vasorum* L1 found in the faeces. One immature *A. vasorum* female extracted from the right eye. | Levamisole injections 10/20 mg/kg for 3 days | [30] |
| 3-year-old; female; Cavalier King Charles Spaniel | UK | Otitis interna, head tilt, submaxillary lymph node enlargement. Presence of a motile worm in the anterior chamber of the eye on a one-day follow up visit. The dog died from a sudden attack of acute respiratory distress | Numerous *A. vasorum* adults in the right ventricle and pulmonary artery and one adult in the anterior chamber detected at post-mortem examination | n/a | [29] |
| 2-year-old; female; Staffordshire Bull Terrier | UK | Post-inflammatory retinal degeneration, episcleral hyperaemia and vitreous haemorrhage. Presence of a motile worm in the anterior chamber of the left eye | *A. vasorum* L1 found in the faeces. No morphological or molecular identification of the nematode found in the left eye | Fenbendazole 50 mg/kg for 10 days | [18] |
| 20-month-old; male; Miniature Dachshund | Canada | Chronic diarrhoea and coughing. Ataxia, depression, and impaired vision. Multiple retinal haemorrhages, iris congestion. The dog was euthanised due to the progression of nervous and ocular diseases | Numerous *A. vasorum* adults recovered from pulmonary artery and disseminated larval infection including both eyes, kidneys, brain, spinal cord, bronchial lymph nodes, heart, intestine and pancreas, detected at post-mortem examination | n/a | [25] |
Additional file

Additional file 1: Case 2. Angiostrongylus vasorum in the anterior chamber of the left eye of a 2-year-old male Cavalier King Charles Spaniel dog. (MOV 1003 kb)

Competing interests
The authors declare that they have no competing interests.

Authors' contributions
VC and DO conceived the study, contributed with data analysis and wrote the first draft of the manuscript. RPL performed the morphological identification. JP, PG, MC, and NdA collected samples and were responsible for all procedures on animals. MSL performed the molecular analysis of samples. DW critically reviewed the manuscript. All authors read and approved the final version of the manuscript.

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References
1. Spratt DM. Species of Angiostrongylus (Nematoda: Metastrongyloidea) in wildlife: A review. Int J Parasitol Parasites Wildl. 2015;4:78–89.
2. Anderson RC. Nematode parasites of vertebrates their development and transmission. 2nd ed. Wallingford: CABI Publishing. 2000.
3. Grewal PS, Grewal SK, Tan L, Adams BJ. Parasitism of molluscs by nematodes: types of associations and evolutionary trends. J Nematol. 2003;35:146–56.
4. Wang QP, Wu ZD, Wei J, Owen RL, Lun ZR. Human Angiostrongylus cantonensis: an update. Eur J Clin Microbiol Infect Dis. 2012;31:389–95.
5. Kim JR, Hayes KA, Yeung RV, Cowie RH. Diverse gastropod hosts of the human Angiostrongylus cantonensis, the rat lungworm, globally and with a focus on the Hawaiian Islands. PLoS One. 2014;9:e89609.
6. Romero-Alegria A, Belhassen-Garcia M, Vefasco-Tirado V, Garcia-Mingo A, Alvela-Suarez L, Pardo-Lledias J, et al. Angiostrongylus costaricensis: systematic review of case reports. Adv Infect Dis. 2014;36:41.
7. Alfaro-Alarcón A, Venzonio V, Galiero G, Cerone A, Gutierrez N, Chinchilla A, et al. First report of a naturally patent infection of Angiostrongylus costaricensis in a dog. Vet Parasitol. 2015;212:3-4:431–4.
8. Morgan ER, Shaw SE, Brennan SF, De Waal TD, Jones BR, Mulcahy G. Angiostrongylus vasorum: a real heartbreaker. Trends Parasitol. 2005;21:49–51.
9. Koch J, Willesea JL. Canine pulmonary angiostrongylosis: an update. Vet J. 2009;179:346–59.
10. Taylor CS, Garcia Gato R, Learmonth J, Attz NA, Montgomery C, Rose H, et al. Increased prevalence and geographic spread of the cardiopulmonary nematode Angiostrongylus vasorum infection in fox populations in Great Britain. Parasitology. 2015;142:1190–5.
11. Elsheikh HA, Holmes SA, Wright I, Morgan ER, Lacher DW. Recent advances in the epidemiology, clinical and diagnostic features, and control of canine cardio-pulmonary angiostrongylosis. Vet Res. 2014;45:92.
12. Di Cesare A, Traversa D, Manzocchi S, Meloni S, Grilloti E, Auriemma E, et al. Elusive Angiostrongylus vasorum infections. Parasit Vectors. 2015;8:438.
13. Colella V, Giannelli A, Bianti E, Ramos RA, Cantacessi C, Dantas-Torres F, et al. Feline lungworms unlock a novel mode of parasite transmission. Sci Rep. 2015;5:13105.
14. Rosen L, Ash LR, Wallace GD. Life history of the canine lungworm Angiostrongylus vasorum (Baillet). Am J Vet Res. 1970;31:131–43.
15. Guilhon J. Larval development of Angiostrongylus vasorum (Baillet, 1866) in the Anoiodae organism. C R Acad Sci Hebd Seances Acad Sci D. 1965;261:4225–7.
16. Bolt G, Monrad J, Frandsen F, Henriksen P, Dietz HH. The common frog (Rana temporaria) as a potential paratenic and intermediate host for Angiostrongylus vasorum. Parasitol Res. 1993;79:428–30.
17. Barcante TA, Barcante JM, Dias SR, Lima WS. Angiostrongylus vasorum (Baillet, 1866) Kamensky, 1905: emergence of third-stage larvae from infected Biomphalaria glabrata snails. Parasitol Res. 2003;91:471–5.
18. Manning SP. Ocular examination in the diagnosis of angiostrongylosis in dogs. Vet Rec. 2007;160:252–5.
19. Lia RP, Traversa D, D’Anna N, Giannelli A, Dantas Torres F, Otranto D. Aberrant ocular infection by Angiostrongylus vasorum in a dog. Roma XXVII Congresso nazionale della società italiana di parasitologia SoPa; 2014.
20. Costa CJ, Costa HM, Guimaraes MP. Redescription of Angiostrongylus vasorum (Baillet, 1866) and systematic revision of species assigned to the genera Angiostrongylus Kamensky, 1905 and Angiostrongylus Schultz, 1951. Rev Med Vet. 2003;154:9–16.
21. McCarty JW, Morgan ER. Identification of first-stage larvae of metastrongyle parasites from dogs. Vet Rec. 2009;165:258–61.
22. Oliveira-Júnior SD, Barcante JMP, Barcante TA, Ribeiro VM, Lima WS. Ectopic localization of a dog with an infected Angiostrongylus vasorum larvae in the eye of a infected dog. Parasitol Res. 2004;121:293–6.
23. Ferdushy T, Hasan MT. Survival of first stage larvae (L1) of Angiostrongylus vasorum under various conditions of temperature and humidity. Parasitol Res. 2010;107:1323–7.
24. Cury MC, Lima WS. Rupture of femoral artery in a dog infected with Angiostrongylus vasorum. Vet Parazitol. 1996;65:313–5.
25. Perry AW, Hering R, Kennedy MJ. Angiostrongylosis with disseminated larval infection associated with signs of ocular and nervous disease in an imported dog. Can Vet J. 1991;32:430–1.
26. Rallat A, Henry A. Contribution a l’étude des nematodes parasites de l’œil du chien. Bul Soc Centr Med Vet Paris. 1913;67:209–15.
27. Henry A, Lebouyres G. Strongle des vaisseaux dans l’œil d’un chien. Bul Soc Centr Med Vet Paris. 1927;80:263–5.
28. Payen D. Angiostrongylus vasorum at a pre-adult phase in the anterior chamber of a young dog’s eye. Munich International Veterinary Ophthalmology Meeting. 2004. p. 125.
29. King MCA, Grose RMR, Startup G. Angiostrongylus vasorum in the anterior chamber of a dog’s eye. J Small Anim Prac. 1994;35:326–8.
30. Rosenlund P, Boserup F, Monrad J. Angiostrongylus vasorum in the anterior chamber of the eye. Euro J Compan Anim Pract. 1995;3:31–3.
31. Weiner DJ, Agurre G, Dubielzig R. Ectopic-site filariid infection with immunologic follow-up of the host. Dallas: Proceedings of the Heartworm Symposium; 1980. p. 51–4.
32. Dantas-Torres F, Lia RP, Barbuto M, Casiraghi M, Crouse A, Caligiani L, et al. Ocular dicrofilariosis by Dicrofilaria immitis in a dog: first case report from Europe. J Small Anim Pract. 2009;50:667–9.
33. Zarfofs MK, Dubielzig RR, Eberhard ML, Schmidt KS. Canine ocular onchocerciasis in the United States: two new cases and a review of the literature. Vet Ophthalmol. 2005;8:51–7.
34. Otranto D, Dantas-Torres F, Giannelli A, Latora MM, Papadopoulos E, Cardoso L, et al. Zoonotic Onchocerca lupi infection in dogs, Greece and Portugal, 2011–2012. Emerg Infect Dis. 2013;19:2000–3.
35. Komenou AT, Thomas AL, Papadopoulos E, Otranto D, Eberhard ML. Intravitreal localization of Onchocerca lupi adult worm in a dog with anterior uveitis: A case report. Vet Ophthalmol. 2015;18:127–30.
36. Schnyder M, Stebler K, Naucke TJ, Lorentz S, Deplazes P. Evaluation of a noninvasive diagnostic method for the detection of Onchocerca lupi in dogs. Int J Parasitol. 2005;35:177–82.
40. Schnyder M, Fahrion A, Ossent P, Kohler L, Webster P, Heine J, et al. Larvicidal effect of imidacloprid/moxidectin spot-on solution in dogs experimentally inoculated with Angiostrongylus vasorum. Vet Parasitol. 2009; 166:326–32.

41. Conboy G, Schenker R, Strehlau G. Efficacy of Milbemax (milbemycin/praziquantel) for the treatment and prevention of Angiostrongylus vasorum infection in dogs. Philadelphia: Proceedings of the Joint 49th Annual Meeting of the American Association of Veterinary Parasitologists/79th Meeting of the American Society of Parasitologists; 2004. p. 92.

42. Punyagupta S, Juttijudata P, Bunag T. Eosinophilic meningitis in Thailand. Clinical studies of 484 typical cases probably caused by Angiostrongylus cantonensis. Am J Trop Med Hyg. 1975;24:921–31.

43. Diao Z, Wang J, Qi H, Li X, Zheng X, Yin C. Human ocular angiostrongyliasis: a literature review. Trop Doct. 2011;41:76–8.