SURVEY OF LUNGWORM INFECTION OF DOMESTIC CATS IN HUNGARY

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From 61 settlements of 12 Hungarian counties, 303 domestic cats were included in this survey. Between autumn 2016 and spring 2018, fresh faecal samples were randomly collected and examined by flotation and by the Baermann–Wetzel method for the presence of lungworm infection. No eggs of *Eucoleus aerophilus* were detected. Morphological identification of first instar larvae (L1) was also carried out. In the faeces of 60 cats (19.8%) from 17 settlements and Budapest, L1 of *Aelurostrongylus abstrusus* were found. More than half of the cats were from the western part of the country. The average number of larvae per gram of faeces was 190.2 ± 304.88. These results are in line with the former findings on the prevalence of aelurostrongylosis of domestic cats in Hungary. In addition, *Oslerus rostratus* was also found for the first time in the faecal samples of three cats from the eastern part of the country, infected also with *Ae. abstrusus*. The average age (2.51 ± 1.26 years) of infected cats indicates that lungworm infection is more common among younger cats. No relationship was found between the lungworm infection and the sex of cats. Non-neutered cats had a significantly higher proportion of lungworm infections. Two-thirds of the infected cats were apparently healthy, and only 19 individuals showed clinical signs of respiratory disorders.

Key words: Lungworms, *Aelurostrongylus abstrusus*, *Oslerus rostratus*, cat, survey, Hungary

Lungworm infection of domestic and wild cats occurs in many countries around the world. It is most commonly caused by *Aelurostrongylus abstrusus*, Railliet, 1898 (Strongylida, Angiostrongylidae), called the cat lungworm (Anderson, 2000; Bowman, 2000). The 5- to 10-mm-long adults reside in the alveolar duct and the terminal bronchioles of the lungs. Over the past decade, other lungworm species of domestic cats such as *Troglostrongylus brevior* and *T. subcrenatus* (Strongylida, Crenosomatidae), *Oslerus rostratus* (syn. *Anafilaroides rostratus*) (Strongylida, Filaroididae) and *Eucoleus aerophilus* (syn. *Capillaria aerophila*) (Enoplida, Trichuridae) have also been reported (Traversa et al., 2009; Jefféries et al., 2010; Brianti et al., 2012, 2014; Di Cesare et al., 2014; Giannelli

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et al., 2017). There have been several cases when two or more lungworm species occurred concurrently in cats (Juste et al., 1992; Jefferies et al., 2010; Traversa et al., 2014; Daikou et al., 2015; Varcasia et al., 2015). With the exception of *E. aerophilus*, the other species develop indirectly with gastropod intermediate hosts. Cats can be infected by intermediate hosts or various other animals (e.g. small mammals, lizards, frogs) which consumed slugs and snails acting as paratenic hosts (Anderson, 2000; Bowman, 2000; Pennisi et al., 2015). Depending on the parasite species, the degree of infection and the immunological status of the host, the parasitosis can be asymptomatic or the cats may show mild to severe clinical signs (coughing, sneezing, tachypnoea). Heavy infection can even be fatal (Traversa et al., 2010; Traversa and Di Cesare, 2013; Philbey et al., 2014; Pennisi et al., 2015).

Data on the distribution and prevalence of cat lungworm infection in Hungary are scarce. In a postmortem examination of 57 stray cats originating from 9 hunting areas of the country *C. aerophila* worms were found in 14 cadavers (Takács and Takács, 2002), *Ae. abstrusus* was detected by autopsy (Kávai, 1977; Dobos-Kovács, 1981; Takács and Takács, 2002) or faecal examination (Capári et al., 2013).

The objectives of this investigation were to enhance the knowledge on the distribution and risk factors of lungworm infection of domestic cats in Hungary.

**Materials and methods**

**Sampling**

Altogether 303 cats from 61 settlements of 12 counties (Table 1 and Fig. 1) were randomly included in the survey. Most cats (*n* = 269) had owners, while the remainder were kept in temporary accommodation. Between autumn 2016 and spring 2018 fresh faecal samples were collected once. Sampling was done with the owners’ permission and assistance.

The approx. 5-gram samples per cat stored in plastic jars having individual numbers were usually taken to the laboratory on the same day or 2–3 days later after storing them in a fridge. Simultaneously, a questionnaire was completed for each cat (e.g. breed, age, sex, keeping, presence/absence of respiratory signs, last anthelmintic treatment).

**Test method**

The *E. aerophilus* eggs were examined by the flotation method using zinc sulphate solution. The lungworm larvae (L1) were isolated by Baermann–Wetzel method (Giannelli et al., 2015). One gram of faecal sample packed in a 10 × 10 cm double-layered gauze was kept in a glass filled with lukewarm water for 24 h.
Table 1

The number of cats sampled per site (N), the number (N) of cats infected with *Aelurostrongylus abstrusus* and *Oslerus rostratus* and the number of infected cats with respiratory signs (e.g. frequent sneezing, running nose)

| Site                     | N  | Ae. abstrusus (N) | O. rostratus (N) | Respiratory signs (N) |
|--------------------------|----|-------------------|------------------|-----------------------|
| Alsózsolca               | 1  | 0                 | 0                | 0                     |
| Badacsony tornaj         | 1  | 1                 | 0                | 1                     |
| Balatonalmádi           | 5  | 0                 | 0                | 0                     |
| Bedő                     | 6  | 6                 | 0                | 4                     |
| Békés                    | 3  | 0                 | 0                | 0                     |
| Békésesaba               | 7  | 0                 | 0                | 0                     |
| Bőcs                     | 2  | 0                 | 0                | 0                     |
| Budaörs                  | 3  | 1                 | 0                | 0                     |
| Budapest                 | 40 | 1                 | 0                | 0                     |
| Csatár                   | 2  | 0                 | 0                | 0                     |
| Csopak                   | 4  | 0                 | 0                | 0                     |
| Debrecen                 | 13 | 0                 | 0                | 0                     |
| Debrecen-Józsa           | 3  | 1                 | 0                | 1                     |
| Dióső                     | 1  | 0                 | 0                | 0                     |
| Dőmösöd                  | 5  | 0                 | 0                | 0                     |
| Dunakeszi                | 2  | 0                 | 0                | 0                     |
| Érd                      | 13 | 2                 | 0                | 0                     |
| Erdőbenye                | 4  | 0                 | 0                | 0                     |
| Gesztely                 | 3  | 0                 | 0                | 0                     |
| Gödöllő                  | 4  | 0                 | 0                | 0                     |
| Gyula                    | 3  | 0                 | 0                | 0                     |
| Hajdúböszörmény           | 17 | 2                 | 2                | 0                     |
| Hajdúdorog                | 9  | 0                 | 0                | 0                     |
| Hajdúzobozdó               | 1  | 0                 | 0                | 0                     |
| Herceghalom              | 1  | 0                 | 0                | 0                     |
| Hernádik                 | 6  | 0                 | 0                | 0                     |
| Hernádikak-Belegrád      | 9  | 7                 | 1                | 6                     |
| Hernádnémeti             | 2  | 0                 | 0                | 0                     |
| Hódmezővásárhely         | 4  | 0                 | 0                | 0                     |
| Ikény                    | 1  | 0                 | 0                | 0                     |
| Ivánca                   | 1  | 1                 | 0                | 0                     |
| Kazincbarcika            | 2  | 0                 | 0                | 0                     |
| Kisalu                   | 1  | 0                 | 0                | 0                     |
| Komárom                  | 1  | 0                 | 0                | 0                     |
| Körösötvég               | 3  | 0                 | 0                | 0                     |
| Mezőberény               | 2  | 0                 | 0                | 0                     |
| Miskolc                  | 8  | 3                 | 0                | 1                     |
| Nyíregyháza              | 6  | 0                 | 0                | 0                     |
| Ólásd                    | 13 | 13                | 0                | 0                     |
| Ökény                    | 1  | 0                 | 0                | 0                     |
| Orosháza                 | 3  | 0                 | 0                | 0                     |
| Sárkapatak               | 6  | 0                 | 0                | 0                     |
| Sátoraljaújhely           | 4  | 0                 | 0                | 0                     |
| Sümeg                    | 7  | 6                 | 0                | 3                     |
| Sümegécsehi              | 2  | 2                 | 0                | 0                     |
| Szeged                   | 10 | 0                 | 0                | 0                     |
| Szentendre               | 3  | 0                 | 0                | 0                     |
| Szentimrefalva           | 8  | 6                 | 0                | 2                     |
| Toceva                   | 2  | 0                 | 0                | 0                     |
| Tura                     | 1  | 0                 | 0                | 0                     |
| Ukk                      | 4  | 4                 | 0                | 1                     |
| Vác                      | 7  | 0                 | 0                | 0                     |
| Vámosszabadi             | 1  | 0                 | 0                | 0                     |
| Vezénző                  | 4  | 1                 | 0                | 0                     |
| Vilmány                  | 3  | 0                 | 0                | 0                     |
| Zalaegerszeg             | 10 | 0                 | 0                | 0                     |
| Zalagymáros              | 2  | 2                 | 0                | 0                     |
| Zalaszegvár              | 1  | 1                 | 0                | 0                     |
| Zalaszentiván           | 4  | 0                 | 0                | 0                     |
| Zalaszentloóne           | 2  | 0                 | 0                | 0                     |
| Zamárdi                  | 4  | 0                 | 0                | 0                     |
After 5 min of centrifugation of the liquid at 600 rpm, the morphological identification of larvae found in the sediment was carried out based on the published descriptions (Traversa and Di Cesare, 2016). The body length and the morphology of head and tail end were considered (Table 2). The identification of *O. rostratus* L1 was confirmed with PCR by Alessio Giannelli in Bari, Italy (personal information). The number of larvae per one gram of faeces (LPG) was determined.

![Map of Hungary with sampling sites](image)

**Fig. 1.** Negative sampling sites (grey dots), sites of *Aelurostrongylus abstrusus* (red dots) and cats dually infected with *Ae. abstrusus* and *Oslerus rostratus* (red triangles)

**Statistical evaluation**

The independence of infection from age group, sex, and keeping mode was analysed by Fisher’s exact test. We used propensity score-based pairing to form sample pairs (Dinya and Solymosi, 2016). A limit value of $P < 0.05$ was used to evaluate the results. All analyses were performed in R environment (R Core Team, 2019).

**Results**

No *E. aerophilus* eggs occurred in the samples. There were *Ae. abstrusus* L1 in the faeces of 60 cats (19.8%; 95% CI: 15.71–24.65) living in 17 settlements and one kept in Budapest (Table 1 and Fig. 1). The LPG values showed significant differences, with an average number of 190.2 ±
304.9. *Oslerus rostratus* L1 were also found in the faecal samples of three cats infected with *Ae. abstrusus*. All infected cats belonged to the European shorthaired breed, most of them (47/60, 78.33%; 95% CI: 66.38–86.87) lived outdoors, and 11 (18.33%; 95% CI: 10.56–29.92) stayed both indoors and outdoors. Only two out of 74 animals kept indoors had lungworm larvae in their samples (2.7% 95% CI: 0.74–9.33). Significantly more animals which lived outdoors or both indoors and outdoors were infected than those kept exclusively indoors (OR: 12.15, 95% CI: 3.07–105.49, P < 0.001). The average age of the infected cats was 2.51 ± 1.26 years, the youngest and the oldest was 8 months and 6 years old, respectively. There was no significant difference (P = 0.5916) between the infection rate of cats younger than one year of age and 1- to 5-year-old cats. The animals under one year and between 1 and 5 years old had a higher risk of lungworm infection than cats older than 5 years (OR: 7.85, 95% CI: 1.93–69.17, P < 0.001 and OR: 6.16, 95% CI: 1.24–60.20, P = 0.0157). The proportion of infected males (34/178, 19.1%, 95% CI: 14.0–25.5) and females (26/125, 20.8%, 95% CI: 14.61–28.73) did not differ significantly. A significantly higher proportion of non-neutered cats were infected with lungworms (OR: 2.98, 95% CI: 1.49–6.32, P < 0.001). When propensity score-based matching was applied, in order to have a balanced population (n = 120) for free range and neutering, the effect of young age remained (OR: 3.83, 95% CI: 1.09–17.22, P = 0.034). Nearly one-third of the infected animals (19/60, 31.67%, 95% CI: 21.31–44.23) showed respiratory signs (e.g. frequent sneezing, running nose) but their number did not differ significantly (P > 0.05) from those of infected cats not showing clinical signs.

**Discussion**

The results of this study are in line with previous findings showing that *Ae. abstrusus* infection of domestic cats is fairly prevalent in Hungary (Kávai, 1977; Dobos-Kovács, 1981; Takács and Takács, 2002; Capári et al., 2013). The occurrence of aelurostrongylosis in new regions can be considered a new finding. In this survey the prevalence of this parasitosis was 19.8%. In the previous pathological or faecal examinations a lower rate of infections was detected: *Ae. abstrusus* worms were found only in 2 out of 50 cats (4%) (Kávai, 1977) and in one out of 57 stray cats (1.7%) (Takács and Takács, 2002) at autopsy. In a parasitological survey of 235 domestic cats carried out in the western part of Hungary, the prevalence of aelurostrongylosis was 14.5% (Capári et al., 2013). In the present study, more than half of the infected animals (n = 35/60, 58.3%) lived in that region. Further studies could answer the question whether the endemic occurrence of this parasitosis in that region is related to environmental and weather conditions more favourable for the intermediate hosts, which are also assumed in other parasitic infections (Patz et al., 2000; Morgan et al., 2009). Lungworm in-
Infections of cats mostly caused by *Aelurostrongylus abstrusus* have been found in several European countries where different prevalence values were reported, e.g. Italy: 8.1–17.8% (Traversa et al., 2008; Di Cesare et al., 2015; Giannelli et al., 2015, 2017), Portugal: 6.25–17.4% (Payo-Puente et al., 2008; Waap et al., 2014; Giannelli et al., 2017), Spain: 1.7–5% (Miró et al., 2004; Giannelli et al., 2017); Denmark: 8.86–13.6% (Olsen et al., 2015; Hansen et al., 2017). By examining more faecal samples collected during 2–3 consecutive days, it may have been possible to find a higher prevalence of lungworm infection than obtained in our study, as the intermittent discharge of larvae was observed in chronically infected cats (Ribeiro and Lima, 2001; Payo-Puente et al., 2008). Recent European studies have reported that at least one of 10 cats is exposed to lungworm infections, the incidence of this parasitosis being more common on the continent than previously thought. In this context, it was noted that lungworm infection of cats is not regarded as a growing threat because there are no previous data to be used for comparison (Giannelli et al., 2017). The authors assumed that the lower occurrence of infected cats found in the western part of Europe can be explained by the more frequent application of anthelmintic drugs.

| Species | Length (μm) | Head | Tail |
|---------|-------------|------|------|
| *Aelurostrongylus abstrusus* | 360–410 | rounded, terminal oral opening | kinked (S shaped), distinct knob-like or small finger-like projections |
| *Troglostrongylus brevior* | 300–360 | pointed, subterminal oral opening | gradually tapered to the extremity, deep dorsal and a shallower ventral incisure |
| *Troglostrongylus subcrenatus* | 270–300 | pointed, subterminal oral opening | gradually tapered to the extremity, deep dorsal and a shallower ventral incisure |
| *Oslerus rostratus* | 330–410 | central oral opening, surrounded by a cuticular ring with dorsal and ventral prominences | with a constriction anterior to the end and tip with a kinked appearance |

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This is a first record for Hungary that in the present study *O. rostratus* was also found in three cats (in the form of co-infection with *Ae. abstrusus*). The 30- to 40-mm-long adults of *O. rostratus* were mostly found in the bronchial submucosa and peribronchial tissues of wild cats and lynx (Brianti et al., 2014a). Since the first description of this nematode species (Gerichter, 1949), it has also been found in domestic cats of some European countries, which may have been infected accidentally (Juste et al., 1992; Millán and Casanova, 2009; Brianti et al., 2014a; Varcasia et al., 2015; Giannelli et al., 2017). Further studies could reveal how three cats kept in two settlements in the eastern part of the country became infected. One of the possible reasons for the rare occurrence of *O. rostratus* in domestic cats is that the living space of wild cats is narrowing, and their number is constantly decreasing in the country (Biró et al., 2009). Except for two infected cats the third one lived outdoors or both indoors and outdoors. A similar result has been reported by others, suggesting that the living conditions have a decisive influence on the infection of animals, since the consumption of naturally infected intermediate or paratenic hosts are necessary for lungworm infections (Genchi et al., 2014; Giannelli et al., 2017). The two cats that were living indoors had probably become infected before they were switched to indoor keeping.

The average age of infected cats (2.51 ± 1.26 years) indicates that lungworm infection is more common among younger cats. Examining the infection rate of 303 animals divided into three age groups, there was a significantly lower risk of infection among animals above 5 years of age due to the acquired immunity (Pennisi et al., 2015). In other studies, there was no difference between the infection rates of the different age groups (Genchi et al., 2014; Tamponi et al., 2014). Some authors have reported that there is a higher risk of infection among young animals (Traversa et al., 2010; Barutzki and Schaper, 2013; Di Cesare et al., 2013). According to other authors, lungworms are more common among adult cats (Mircean et al., 2010; Knaus et al., 2014). We agree with those researchers who say that cats can be infected with lungworms at any age if they can reach the infective intermediate or paratenic hosts in the environment (Beugnet et al., 2014; Giannelli et al., 2015).

No relationship was found between lungworm infection and the sex of animals, as the parasitosis occurred in almost the same proportion in males and females, as opposed to other findings (Traversa et al., 2008; Tamponi et al., 2014). We agree with Traversa et al. (2008) and Genchi et al. (2014) who reported that the sex of cats did not affect their infection rate. Non-neutered cats had a significantly higher proportion of lungworm infections, which is probably due to a decrease in their activity after the spaying operation.

Two-thirds of the infected cats were free of clinical signs, and only 19 showed respiratory signs. Italian researchers reported that more than 50% of animals infected with *Ae. abstrusus* had clinical signs (Traversa et al., 2008; Genchi et al., 2014; Di Cesare et al., 2015). The question arises whether or not there
is a correlation between LPG values and the occurrence of clinical signs. Contradic-
tory data have been reported so far. There were scientists who did not find
that LPG values were significantly higher in cats with clinical signs (Traversa
and Di Cesare, 2016). Others have reported that the development of clinical signs
results from the massive egg production of worms and the lesions produced in
the lungs mainly by L1. Therefore, higher levels of LPG are found in such indi-
viduals (Naylor et al., 1984; Gerdin et al., 2011; Genchi et al., 2014).

The knowledge regarding lungworm infection of domestic cats living in
Hungary has been expanded with the results of the current studies. However, the
following questions have arisen, which need to be answered by further studies:
What is the frequency of the parasitosis caused by *O. rostratus* in domestic cats?
Do *Troglostrongylus* spp. occur in the country? What are the most common in-
termediate hosts of the nationwide distributed *Ae. abstrusus*? Can the geograph-
ical distribution of cat lungworms be influenced by the climate change in the
country?

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