

HELMINTHOLOGIA, 54, 3: 199 - 210, 2017

Segregated settlements present an increased risk for the parasite infections spread in Northeastern Slovakia

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Article info

Received March 14, 2017
Accepted June 7, 2017

Summary

The occurrence of parasitic infections among the children, dogs and its association with soil contamination in two villages with different hygiene level standards were analysed. Infections were present in both examined localities, but in the village with higher living standard, a better personal and communal hygiene level and better dogs care a lower occurrence of parasitic germs in soil was detected. High prevalence of protozoa and helminths was observed not only within canine population but also in children throughout the year in the village with lower hygiene and socio-economic standard. We have identified up to 12 taxa of parasites in 127 collected dogs' excrements and mean prevalence was 71.65 %. The most frequent were eggs of family Ancylostomatidae and *Ascaris* spp., followed by *Toxocara canis*, *Toxascaris leonina*, *Giardia duodenalis* cysts, *Isospora* spp. oocysts, eggs of *Capillaria aerophila*, *Trichuris vulpis*, *Taenia* type eggs, *Dipylidium caninum*, oocysts of *Sarcocystis* spp. and larvae of *Angiostrongylus vasorum*. The soil samples collected near dwellings were highly contaminated. Two thirds of samples contained eggs for the most part of family Ancylostomatidae as well as genera *Ascaris* and *Toxocara*. Among the kids population helminth ova were present in 53.17 % of stool samples, where the eggs of *Ascaris lumbricoides*, *Trichuris trichiura*, *Enterobius vermicularis*, *Hymenolepis diminuta* and cysts of *G. duodenalis* were the most frequent. In contrast, parasitic diseases were not seen in children population living in the locality with common hygiene standard.

Keywords: protozoa infection; helminth infection; soil; dogs; children; Northeastern Slovakia

Introduction

Parasitic infections present still a serious problem of the 21st century. The WHO Secretariat (2012), Røttingen *et al.* (2013) and von Philipsborn *et al.* (2015) classified diseases caused by helminths, roundworms, hookworms, whipworms and tapeworms as type III diseases. It means that above mentioned diseases are noticeably linked with poverty level and socio-economic status of its population. Elyana *et al.* (2016) reported that intestinal parasitic infections are major public health problem. Particularly in danger are the children living in poor or rural neighbourhoods. Globally, about

1.5 billion people are infected with at least one intestinal parasitic species (Pulan *et al.*, 2014; Hotez *et al.*, 2014). Pulan *et al.* (2014) reported that in 2010 about 438.9 million people were infected with hookworms. Additionally, 819.0 million *Ascaris lumbricoides* and 464.6 million *Trichuris trichiura* infections were identified.

Among of a many parasitic infections, we should principally mention those which are transmitted through the soil or water. They are represented by endoparasites such as *Giardia duodenalis*, *Cryptosporidium* spp., *Trichuris* spp., *Ascaris* spp., *Ancylostoma* spp., *Toxocara* spp., *Echinococcus* spp., and *Toxascaris* spp. (Bajer, 2008; Traversa *et al.*, 2014). For diseases caused by endopara-

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sites the most likely route of human infection is fecal-oral transmission followed by the contact with infected humans and animals (wild, stray, domestic), with contaminated food, soil, water, or environment. The main source of the infection is environment contaminated with feces from infected animals living in close vicinity with the man (Despomer, 2003; Szabová *et al.*, 2007; Macpherson, 2013; Ondriska *et al.*, 2013). Parasitic infections are very often transmitted at places with high population density, poor hygiene conditions and low socioeconomic status of community (Adeoye, *et al.*, 2007; Rudohradská *et al.*, 2012; Papajová *et al.*, 2014).

The prevalence of intestinal parasitic diseases in Slovakia is due to its geographical location and good hygiene conditions quite low. However, it may be easily transmitted by socially disadvantaged people living in a low hygiene conditions. Primarily these diseases occur in the population of marginalised communities, which are distinct by complete social exclusion, due to a various factors. In Slovakia, according to the performed research and governmentally released strategy papers the groups put most at risk are the Roma people. This is in Slovakia a very specific problem (Madarasová Gecková *et al.*, 2014). According to the Atlas of Roma communities (Mušinka *et al.*, 2014) there are 803 Roma settlements in Slovakia territory. It is home of more than 215 000 people living inside the villages (30.7 %), at the outskirts of the villages (40.4 %) or in segregated settlements (16.6 %). Most of the Roma population lives with the major population (46.5 %) (Filadelfiová *et al.*, 2007; Mušinka *et al.*, 2014). Roma ethnic group inhabit shacks and houses in the villages, apartments with reduced living standards and apartments found on abandoned houses in urban areas (Dubayová, 2001). In the cities, we see urban areas which are exclusively inhabited by Roma people commonly called – Roma ghettos (Mušinka, 2002).

This study focuses on the analysis of the potential health risks caused by parasites in rural locations of Slovakia. The aim of the study was to monitor the occurrence of intestinal parasites in dogs in two Northeastern Slovakian localities with different hygiene standards. The soil contamination near dwellings was evaluated for the period of one year. A coprological diagnostics of intestinal parasites in the children population was also performed.

Material and Methods

Characterization of the studied localities

This study was performed in two neighbouring villages in Prešov County (Northeastern Slovakia) that wished to remain anonymous and are further specified as “village A” (about 500 inhabitants) and “village B” (about 510 inhabitants). Both these locations are rural, but differ by hygiene standards, levels of sanitary conditions and socio-economic status of dog owners. The village A is inhabited predominantly by Roma minority group (more than 95 %) and characterized by a low level of environmental hygiene. Electricity is available for public lighting and to some houses. Clean water

supply and sewage systems are not available. Water for the residents is available only through water wells. There are 40 registered dogs which are either tied by chain to the kennel, or moving freely around settlements. They are usually fed on leftovers and households garbage. No data about unregistered animals were available. The neighbouring village B is represented by a higher hygiene and living standards where the gas, electricity, municipal water supply and municipal sewerage are available. Slovak majority population lives in village B where 57 dogs and 10 cats are registered. Dogs are usually kept in the yard and fed by commercially manufactured dry dog food from various manufacturers. Both villages in this study were visited once a month and inspected for the presence of parasites in feces and soil samples.

Coprological examination of excrements

Totally 199 dog fecal samples were collected and examined for the presence of parasite developmental stages. Dog feces have been collected from around the dwellings, public places or taken directly by the owners from backyards. After collection, fecal samples were stored without any conservation at 4 °C and transferred to the laboratory at the Institute of Parasitology SAS in Košice for parasitological examination which was performed within 24 – 48 h. Flotation method with the sucrose flotation solution (specific gravity of 1.27) was used for coprological examination where 3 grams of fecal sample mixed with water was centrifuged for 5 minutes at 1200 rpm (Eppendorf 5804, Germany). After pouring out the supernatant the sucrose flotation solution was added in the test tube. The sediment was than stirred and centrifuged again. After 5 minutes the test tube was refilled with flotation solution (until a meniscus formed) and covered with cover glass. After one hour of egg flotation the coverslip was removed and placed on the glass slide. For the detection of *Giardia* cysts the zinc sulphate flotation solution (specific gravity of 1.18) was used for each fecal sample. All samples were further examined under the light microscope at 20x and 40x magnification (Leica Microsystems, DM 5000B light microscope, Germany).

Parasitological examination of soil samples

In order to identify the presence of parasitic germs in the environment totally 65 soil samples were collected within the vicinity of human settlements and around the kennels and dog pens. 32 samples were from village A and 33 from village B. The sand samples were surveyed according to the Kazacos (1983). Briefly, 100 g of pooled sand sample, 100 ml of water and 0.5 ml of Tween 40 were mixed and decanted for 10 minutes. Subsequently the samples were sieved and replenished with 1000 ml of water. After one hour sedimentation the soil samples were centrifuged (Eppendorf 5804, Germany) and then floated with sucrose flotation solution (specific gravity of 1.3). Samples were examined under the light microscope at 20x and 40x magnification (Leica Microsystems, DM 5000B light microscope, Germany).

Table 1. Prevalence of the propagative parasitic stages in dogs' excrements.

	Negative	Positive	Total	Prevalence (%)	95% CI min	95% CI max	OR	95% CI min	95% CI max	χ^2	P-value
Village A	36	91	127	71.65	63.27	78.76	10.47	5.2	21.08	50.25	<0.0001
Village B	58	14	72	19.44	11.95	30.03					

95% CI – 95% Confidence Interval, OR – Odds Ratio

Table 2. Prevalence of endoparasite taxa detected in dogs' excrements collected from public places.

	Village A (n-127) (%)	Village B (n-72) (%)
<i>Toxocara canis</i>	11.02	8.33
<i>Toxascaris leonina</i>	9.45	8.33
<i>Trichuris vulpis</i>	2.36	1.39
<i>Ascaris</i> spp.	40.94	ND
family Ancylostomatidae	50.39	ND
<i>Capillaria aerophila</i>	3.94	ND
<i>Dipylidium caninum</i>	0.79	ND
<i>Angiostrongylus vasorum</i>	0.79	ND
<i>Taenia</i> type eggs	0.79	ND
<i>Isospora</i> spp.	6.30	ND
<i>Sarcocystis</i> spp.	0.79	2.78
<i>Giardia duodenalis</i>	9.45	ND

n - number of examined samples, ND – not detected

Table 3. Occurrence of the propagative parasitic stages in soil.

	Negative	Positive	Total	Prevalence (%)	95% CI min	95% CI max	OR	95% CI min	95% CI max	χ^2	P-value
Village A	11	21	32	65.63	48.32	79.59	8.59	2.73	27.04	15.06	<0.0001
Village B	27	6	33	18.18	8.61	34.39					

95 % CI – 95 % Confidence Interval, OR – Odds Ratio

Table 4. Occurrence of endoparasite taxa in soil collected from public places.

	Village A (n-32) (%)	Village B (n-33) (%)
<i>Toxocara</i> spp.	34.37	3.03
<i>Toxascaris leonina</i>	3.13	15.15
<i>Trichuris</i> spp.	3.13	ND
<i>Ascaris</i> spp.	50.00	ND
family Ancylostomatidae	43.75	6.06
<i>Capillaria</i> spp.	3.13	ND
<i>Sarcocystis</i> spp.	3.13	ND

n – number of examined samples, ND – not detected

Table 5. Prevalence of propagative parasitic stages in children's stool.

	Negative	Positive	Total	Prevalence (%)	95% CI min	95% CI max	OR	95% CI min	95% CI max	χ^2	P-value
Village A	96	109	205	53.17	46.34	59.88	164.53	10.06	2691.75	63.12	<0,0001
Village B	72	0	72	0	0	5.07					

95 % CI – 95 % Confidence Interval, OR – Odds Ratio

Table 6. Prevalence of endoparasite taxa in children's stool.

	Village A (n=205) (%)	Village B (n=72) (%)
<i>Ascaris lumbricoides</i>	52.20	ND
<i>Trichuris trichiura</i>	2.44	ND
<i>Enterobius vermicularis</i>	0.98	ND
<i>Hymenolepis diminuta</i>	0.49	ND
<i>Giardia duodenalis</i>	8.29	ND

n – number of examined samples, ND – not detected

Parasitological examination of stool samples

Totally, 277 children's stool samples were collected into the plastic containers. After an informed consent was signed by parents or legal guardians stool containers with unique identifiers were handed out together with the instruction how to return them. Containers with stool samples (up to 15g of morning stool) were stored in refrigerator without any conservation at 4 °C and transferred to the laboratory at the Institute of Parasitology SAS for the examination which was performed within 24 – 48 h. Samples were examined with commercially available kit (Paraprep L, Mondial, France). Briefly, for each stool sample 2 ml of ethyl acetate solution and 0.5 g of stool sample was added to 6 ml of 10 % formalin in a mixing chamber. The chamber was then connected through filter with a conical collection chamber. Mixed content was incubated for 24 h at room temperature and the tube was centrifuged at 1000 rpm for 1 minute (Eppendorf 5804, Germany). The entire samples volumes were collected into the collection chambers. The supernatant was discarded and the sediment placed on microscope slides and covered with coverslip. The entire area was examined at 20× and 40× magnification with Leica DM 5000B light microscope (Leica Microsystems, Germany).

Statistical methods

Statistical analysis was performed using Vassar Stats (www.vassarstats.net). 95 % confidence intervals (95 % CIs) and Odds ratio (OR) with 95 % confidence intervals (95 % CIs) were determined. To compare the potential risk factors a Chi-square test (χ^2 ; if all expected cell values were equal to or greater than 5) or a Fisher's exact test (if any expected cell value was less than 5) were used.

Results

Out of 199 examined dogs' fecal samples 52.76 % of them contained parasitic germs. Intestinal parasites infections were detected in both examined villages. There were considerable differences in the prevalence and species diversity of detected helminths between village A and B. Meanwhile the prevalence among dogs from the village A was 71.65 % (95 % CI: 63.27 – 78.76 %) the prevalence of infection among dogs from the village B was only 19.44 % (95 % CI: 11.95 – 30.03 %). This difference was highly significant ($X^2 = 50.25$, $P < 0.0001$) (Table 1). The odds for the infection were higher in dogs from the village A than dogs from the village B (OR = 10.47, 95 % CI: 5.20 – 21.08).

In 127 examined samples from the village A (low hygienic standards) 12 different taxa of parasites were detected. For the most part the eggs of Ancylostomatidae family (50.39 %), *Ascaris* spp. eggs (40.94 %) *Toxocara canis* eggs (11.02 %), eggs of *Toxascaris leonina* (9.45 %), *Giardia duodenalis* cysts (9.45 %), oocysts of *Isospora* spp. (6.30 %), *Capillaria aerophila* eggs (3.94 %), *Trichuris vulpis* eggs (2.36 %), *Taenia* type eggs (0.79 %), *Dipylidium caninum* eggs (0.79 %), *Angiostrongylus vasorum* larvae (0.79 %) and *Sarcocystis* spp. oocysts (0.79 %) were present. In village B

(high hygienic standards) only 4 taxa of parasites in 72 dogs' excrements were found. To be exact the eggs of *T. leonina*, *T. canis*, *T. vulpis* and *Sarcocystis* spp. oocysts were detected (Table 2).

High prevalence of the endoparasitic developmental stages in dogs' feces present a risk factor for the soil contamination, and consequently to the population living in such locality. For this reason, we have examined the occurrence of helminth eggs in the soil. Out of 65 collected soil samples parasites were detected in 27 (41.54 %) of them. The soil samples from the village A were contaminated more than samples from village B. In village A 65.63 % of soil samples contained parasites, while in village B only 18.18 % positivity was observed. This difference was highly significant ($X^2 = 15.06$, $P < 0.0001$, OR = 8.59; Table 3). In a village A, *Ascaris* spp. (50.00 %) and eggs from the family Ancylostomatidae (43.75 %) as well as the *Toxocara* spp. (34.37 %) were the most frequent. As shown on Table 4, the eggs of *T. leonina*, *Trichuris* spp., *Capillaria* spp. and oocysts of *Sarcocystis* spp. occurred occasionally in examined soil samples. The most prevalent parasitic taxa in soil samples from village B were *T. leonina*, *Toxocara* spp. and eggs of the family Ancylostomatidae (Table 4).

Overall, 277 fecal samples of children from village A (205 children) and village B (72 children) were examined for the presence of parasites. The overall infection prevalence in children living in village A was 39.35 % (109/277). No children from the village B were found to be positive for the presence of parasites. In contrast the prevalence of propagative parasitic stages among children from the village A was 53.17 %. This difference was highly significant ($X^2 = 63.12$, $P < 0.0001$) and Roma children had higher odds to become infected by parasites than children living in village B (OR = 164.53, Table 5). Despite living in the close proximity both groups children (Roma and non-Roma) still maintain their specific life style what include different hygiene standards and live with pets. The explanation is obvious. Risk factors in urban settings were not disrupted and both groups of children remained segregated. In children from the village A, the most prevalent parasite was *A. lumbricoides*. Ascarid eggs were found in 52.20 % of the stool samples (107/205). The cysts of *G. duodenalis* were found in 8.29 % samples (17/205) and eggs of *T. trichiura* in 2.44 % (5/205; Table 6). *Hymenolepis diminuta* (0.49 %) eggs were observed less often. However, the perianal swabs and adhesive cellophane-tape tests for the prevalence of *Enterobius vermicularis* were not performed. The eggs of *E. vermicularis* were found unintentionally in the stool (Table 6).

The gastrointestinal parasites infestation in dogs and children persisted throughout the year (Tables 7 – 9). *G. duodenalis* infections in canine and children population in village A were observed particularly during winter and spring (Table 7 and 9). In the village A – location represented by lower hygienic standards, at least one soil sample from monthly collection contained parasitic germs. Most of helminth eggs have developed into larvae or reached various developmental stages. The egg counts per sample varied from 1 to 88 (per 100 g of dry soil). The egg counts were highest in the

Table 7. The monthly occurrence of endoparasites in dogs' excrements collected from public places in village A.

	XI.	XII.	I.	II.	III.	IV.	V.	VI.	VII.	VIII.	IX.	X.
	(n-16)	(n-15)	(n-8)	(n-6)	(n-12)	(n-13)	(n-13)	(n-7)	(n-8)	(n-8)	(n-8)	(n-13)
	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)
<i>Toxocara canis</i>	1	3	4	0	0	1	2	0	0	0	1	2
<i>Toxascaris leonina</i>	0	3	1	0	0	3	2	0	1	0	0	2
<i>Trichuris vulpis</i>	0	0	2	0	0	0	0	0	1	0	0	0
<i>Ascaris</i> spp.	6	7	5	1	6	9	6	1	1	5	4	1
family Ancylostomatidae	9	10	4	3	10	6	5	3	3	3	5	3
<i>Capillaria aerophila</i>	0	0	0	0	0	1	0	2	0	1	1	0
<i>Dipylidium caninum</i>	0	0	0	0	1	0	0	0	0	0	0	0
<i>Angiostrongylus vasorum</i>	0	0	0	0	1	0	0	0	0	0	0	0
<i>Taenia</i> type eggs	1	0	0	0	0	0	0	0	0	0	0	0
<i>Isospora</i> spp.	0	3	1	0	1	0	1	2	0	0	0	0
<i>Sarcocystis</i> spp.	0	0	0	0	0	0	0	1	0	0	0	0
<i>Giardia duodenalis</i>	1	4	0	0	3	0	4	0	0	0	0	0

n – number of examined samples, p – number of positive samples

Table 8. The monthly occurrence of endoparasites in dogs' excrements collected from public places in village B.

	XI.	XII.	I.	II.	III.	IV.	V.	VI.	VII.	VIII.	IX.	X.
	(n-6)	(n-6)	(n-3)	(n-11)	(n-10)	(n-6)	(n-5)	(n-6)	(n-6)	(n-6)	(n-4)	(n-3)
	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)
<i>Toxocara canis</i>	2	2	0	1	0	0	0	0	0	1	0	0
<i>Toxascaris leonina</i>	0	0	1	0	1	0	0	0	0	0	3	2
<i>Trichuris vulpis</i>	0	0	1	0	0	0	0	0	0	0	0	0
<i>Ascaris</i> spp.	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
family Ancylostomatidae	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
<i>Capillaria aerophila</i>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
<i>Dipylidium caninum</i>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
<i>Angiostrongylus vasorum</i>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
<i>Taenia</i> type eggs	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
<i>Isospora</i> spp.	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
<i>Sarcocystis</i> spp.	0	0	0	0	0	0	0	1	0	0	1	0
<i>Giardia duodenalis</i>	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND

n – number of examined samples, p – number of positive samples, ND – not detected

centre of village and the number of eggs was reduced by distance. The most widespread were *Ascaris* spp. eggs which were detected in 16 soil samples. The eggs from family Ancylostomatidae (14 samples) and *Toxocara* spp. eggs (11 samples) were also found quite regularly. In opposite, quite rare was the occurrence of *T. leonina*, *Trichuris* spp. and *Capillaria* spp. (Table 10). In contrast, the soil samples from the village B the developmental parasitic stages were detected in samples collected in January, March, April and September. Only three taxa were identified: *T. leonina* (5 samples), eggs from the family Ancylostomatidae (2 samples) and *Toxocara* spp. (1 sample; Table 11).

Discussion

In 27 countries representing the EU it is estimated that 16 % of its citizens (about 80 million people) live below the poverty level. Poverty, defined as 60% of particular country median income in Europe, is distributed fairly alongside well-defined quartiles. The gross domestic product (GDP) per capita organizes countries into 4 quartiles. Most of the countries in the top two quartiles are represented by western European nations. All the poorest and lowest GDP per capita countries are geographically located in southern and eastern part of the European region. (Hotez & Gurwith, 2011). Intestinal helminth and protozoan infections occur not only among those living in the poverty but also in the more affluent part of Europe. In Slovakia, the population living in poverty is concentrated typically in segregated Roma settlements. Life in the shelters, on the outskirts of towns and villages, or in the forests vicinity expose the Roma people more to the wildlife and the other potential biological agents (mosquitoes, ticks, flies, etc.) thus allowing infections to spread. At the same time, on a small area, a large number of people live together with domestic animals that are often without a proper veterinary control. Moreover, within the vicinity of these settlements animals' excrements and human feces concentrate without an appropriate sanitary control what represents a significant risk for the circulation of soil transmitted infections of animals and man.

The occurrence of intestinal helminth and protozoan infections in children represent a health problem especially in the rural areas with low standard of hygiene (Rudohradská *et al.*, 2012; Harhay *et al.*, 2010; Al-Mekhlafi *et al.*, 2016). Therefore, we determined and compared the occurrence of parasitic infections in dogs, children population and soil found in two neighbouring villages with different levels of hygiene and socio-economic conditions of the population.

Parasitic infections were present in both study areas, but in the village B with a higher standard of living, better personal and communal hygiene levels and better dogs' care a lower occurrence of the parasitic infections in dogs and less contaminated soil were recorded. The total prevalence of intestinal parasites among dogs from the village A with predominance of Roma minority was 71.65 % and 12 different taxa with predominance of family Ancylostomatidae and *Ascaris* spp. were identified. The presence

of *Ascaris* spp. provides evidence dogs coprophagous habits of dogs. Ascariasis spreads primarily by fecal contamination of the environment around human dwellings because of defecation outside the toilets. Such behaviour is called promiscuity defecation (Traub *et al.*, 2002). Such high prevalence numbers bear a resemblance with the situation in developing countries in Asia, Africa or South America. Traub *et al.* (2002) found that 94% of dogs living in poor communities in India eliminate the eggs from the family Ancylostomatidae and 31 % dogs pass the eggs of *Ascaris* spp. In a resource-limited urban community in Gauteng, South Africa, 88 % of dogs have been infected by *Ancylostoma caninum* and 36 % were harbouring *T. canis* (Minnaar & Krecek, 2001). Rubel *et al.* (2003) showed that the higher toxocarosis infection rate was associated with lower financial income. Also hookworm and whipworm infections occur more often in economically deprived localities (Rubel & Wisnivesky, 2005).

The mean prevalence of intestinal parasites in dogs from village with better hygiene was 19.44 % and only 4 taxa were detected. *T. leonina* and *T. canis* appeared to be most frequent. This was followed by *Sarcocystis* spp. and *T. vulpis*. Regarding the dogs fecal samples, *T. leonina* was the most often detected species in the soil samples from village with better environmental hygiene. Previous studies from Slovak Republic show that 45.1 % of dogs were infected by intestinal parasite, especially by *T. canis* (21.9 %), hookworms (18.4 %), whipworms (10.0 %) or *T. leonina* (7.3 %); (Szabová *et al.*, 2007). Our results are comparable to the records from neighbouring countries. Helminth prevalence among dogs in Poland was 34.2 % in rural locations and 56.5 – 80.9 % in dog shelters. The eggs of the nematodes from family Ancylostomatidae were the most frequent, followed by *T. vulpis* and *T. canis*. *T. leonina* was present among dogs in shelters, but was missing among rural dogs (Borecka, 2004). Very similar infection rates were found among dogs from the rural areas in the Czech Republic, where 41.7 % of dogs harboured some species of intestinal parasites. However the most prevalent was *T. canis* and the eggs from family Ancylostomatidae were detected only sporadically (Dubná *et al.*, 2007).

The occurrence of *G. duodenalis* in dogs from the village A with low hygienic standard has also been recorded. Study of Marangi *et al.* (2008) evaluated the presence of giardiasis in dogs and children living in a disadvantaged and socially deprived small Roma community found that 5/14 (35 %) children were found positive by microscopic examination. Additionally 8/14 (57 %) dogs tested both, microscopically and by molecular methods were positive for *G. duodenalis*. In the territory of Eastern Slovakia the giardia infections frequency was examined by Szabová *et al.* (2007). The overall prevalence of the *Giardia* cysts in dogs' excrements from shelters in the cities Trebišov, Košice and Zvolen was 1.6 %. Similarly, Goldová *et al.* (2011) found that the giardiasis prevalence in dogs less than 7 months old was 69.1 %. Meanwhile in older animals the prevalence dropped to 36.9 %. The activities associated with the maintenance and landscaping of urban green spaces may

Table 9. The monthly occurrence of endoparasites in children from the village A.

	XI.	XII.	I.	II.	III.	IV.	V.	VI.	VII.	VIII.	IX.	X.
	(n-18)	(n-20)	(n-15)	(n-10)	(n-18)	(n-20)	(n-27)	(n-16)	(n-20)	(n-10)	(n-11)	(n-20)
	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)
<i>Ascaris lumbricoides</i>	7	10	9	9	11	7	25	6	0	1	5	17
<i>Trichuris trichiura</i>	2	1	0	1	0	0	0	0	0	0	0	1
<i>Enterobius vermicularis</i>	0	0	0	0	1	0	0	0	0	0	0	1
<i>Hymenolepis diminuta</i>	0	0	0	1	0	0	0	0	0	0	0	0
<i>Giardia duodenalis</i>	1	0	0	0	0	0	16	0	0	0	0	0

n - number of examined samples, p - number of positive samples

Table 10. The monthly occurrence of endoparasites in soil from the village A.

	XI.	XII.	I.	II.	III.	IV.	V.	VI.	VII.	VIII.	IX.	X.
	(n-3)	(n-3)	(n-2)	(n-0)	(n-3)	(n-3)	(n-3)	(n-3)	(n-3)	(n-3)	(n-3)	(n-3)
	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)
<i>Toxocara</i> spp.	0	1	2	-	2	1	0	1	1	1	1	1
<i>Toxascaris leonina</i>	0	0	0	-	1	0	0	0	0	0	0	0
<i>Trichuris</i> spp.	0	0	0	-	1	0	0	0	0	0	0	0
<i>Ascaris</i> spp.	1	1	2	-	2	2	0	1	3	2	1	1
family Ancylostomatidae	2	2	2	-	1	1	1	0	1	0	2	2
<i>Capillaria aerophila</i>	0	0	0	-	1	0	0	0	0	0	0	0
<i>Sarcocystis</i> spp.	0	0	0	-	1	0	0	0	0	0	0	0

n - number of examined samples, p - number of positive samples

Table 11. The monthly occurrence of endoparasites in soil from the village B.

	XI.	XII.	I.	II.	III.	IV.	V.	VI.	VII.	VIII.	IX.	X.
	(n-3)	(n-3)	(n-3)	(n-0)	(n-3)	(n-3)	(n-3)	(n-3)	(n-3)	(n-3)	(n-3)	(n-3)
	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)	(p)
<i>Toxocara</i> spp.	0	0	0	-	1	0	0	0	0	0	0	0
<i>Toxascaris leonina</i>	0	0	1	-	2	1	0	0	0	0	1	0
<i>Trichuris</i> spp.	ND	ND	ND	-	ND	ND	ND	ND	ND	ND	ND	ND
<i>Ascaris</i> spp.	ND	ND	ND	-	ND	ND	ND	ND	ND	ND	ND	ND
family Ancylostomatidae	0	0	1	-	1	0	0	0	0	0	0	0
<i>Capillaria aerophila</i>	ND	ND	ND	-	ND	ND	ND	ND	ND	ND	ND	ND
<i>Sarcocystis</i> spp.	ND	ND	ND	-	ND	ND	ND	ND	ND	ND	ND	ND

n - number of examined samples, p - number of positive samples

also contribute to the dispersion of dogs' and cats' excrements on large areas. After the excrements break-down all parasitic propagative stages disperse in the soil. Such contaminated soil become a source of infections and reinfections for animals, and poses a significant risk to man as well. Gawor *et al.* (2008) studied the epidemiological link between the likelihood of reinfection with soil transmitted helminths and children diagnosed with toxocarosis. Examination of the soil from around the homes of infected children have confirmed that more of the *Toxocara* spp. eggs are being found in rural locations than in samples collected from the urban environment. In contrast Antolova *et al.* (2015) studied seroprevalence of human *Toxocara* infections. Cross-sectional study confirmed the 22.1% seropositivity to *Toxocara* in 429 examined Roma inhabitants living in segregated settlements. Meanwhile only 4 of 394 samples from the non-Roma population were found to be positive.

Soil contamination by endoparasites developmental stages in areas inhabited with marginalized groups has been monitored by Rudohradská *et al.* (2011). The settlements environment where children lived and played was contaminated extensively with the parasite propagative stages, where up to 81.6 % of examined soil samples were contaminated with helminths eggs (*Ascaris* spp., *Toxocara* spp., *Trichuris* spp. and strongyloid eggs). Occasionally the eggs of *T. leonina* and *Spirocerca lupi* were detected. The highest soil contamination was detected in villages Jarovnice, Zemplínska Teplica and Sečovce. Research of Štrkolcová *et al.* (2017) in segregated settlement Medzev (Eastern Slovakia) showed that *Strongyloides stercoralis* infections in children and dogs are frequent. Despite similar level of specific antibodies against *S. stercoralis* in all examined dogs Roma children showed higher seroprevalence (33.3 %) when compared with non-Roma children (23.8 %).

However, this study did not investigate the occurrence of cryptosporidiosis – a parasitic disease considered to be a widespread zoonosis. Its occurrence in Roma children was studied by Hasajová *et al.* (2014). Examination of 53 Roma and 53 non-Roma fecal samples led to the conclusion that the risk of *Cryptosporidium* infection is 12 times higher in the Roma children when compared to the non-Roma children. Moreover the occurrence of microsporidia as emerging pathogens in Slovak Roma children and their impact on public health was confirmed (Halánová *et al.*, 2013).

Our results show that the high endoparasitic occurrence in the environment poses a significant risk to the human health. Most exposed are children of preschool age who play in public places what may result in the ingestion of contaminated soil. It should be stated that for example eggs of soil-transmitted helminths in feces need some time to embryonate and become infective because they are not immediately infective (Knop *et al.*, 2012; Grimes *et al.*, 2016). We have detected intestinal parasitic infections in 53.17 % of examined Roma children. The predominant infections found were *A. lumbricoides* and *G. duodenalis* origin. Similarly, Rudohradská *et al.* (2012) studied the occurrence of selected intestinal endoparasites in 81 children living in eastern Slovakia in areas char-

acterized by low hygienic and socioeconomic status. 56.8 % of children were positive for the presence of intestinal parasites. *A. lumbricoides* was found to be the most prominent intestinal parasite (24.7 %). This was followed by *T. trichiura* (17.3 %) and *Taenia* type eggs (4.9 %). *Cryptosporidium* spp. (44.4 %) and *G. duodenalis* (24.7 %) were also detected in a quite high prevalence.

Due to the dogs free movement the soil around houses was highly contaminated with helminths eggs and protozoan cysts and served as a source of the infection spread. Thus, in addition to its own parasite fauna, human intestinal parasites are often passaged through the dogs' digestive system. Triggered by limited availability of sanitary facilities and the absence of sewage system in the village children also participated in soil contamination as well. One of the efficient ways for the reduction of parasitic infections is establishment of a reliable veterinary care. Regular dogs deworming, especially by the owners with small children, reduces the possibility of parasites spread. Introduction of a proper environmental hygiene measures, education in a basic hygiene habits and regular removal of dog excrements will prevent dispersion of parasite developmental stages into the environment. The results of this study were submitted to the local health and social services to utilise data further. Based on our results the preventive measures will be implemented through local community centres and physicians.

Acknowledgement

This work was supported by the scientific Grant Agency of the Ministry of Education of the Slovak Republic, by support from the Slovak Academy of Sciences, VEGA no. 2/0125/17 (0.6). The publication has been realised within the scope of the projects performed in the Centre of Excellence for Parasitology (Code ITMS: 26220120022) based on funding support from the Operational Programme "Research & Development", which is funded by the European Regional Development Fund (0.4).

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