Surveillance of *Echinococcus* tapeworm in coyotes and domestic dogs in Winnipeg, Manitoba

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**Abstract**

**Background:** The *Echinococcus* species, including *E. multilocularis* and *E. canadensis*, are tapeworms that primarily infect canids such as dogs, foxes and coyotes, but which can also infect humans. In humans, *E. multilocularis* can cause alveolar echinococcosis; a serious condition that mimics metastatic malignancy and has a poor prognosis. It is known that coyotes in rural Manitoba are infected with *Echinococcus* species, but it is not known if coyotes in peri-urban areas are also infected.

**Objectives:** To document and map *Echinococcus* species in wild canids and domestic dogs in Winnipeg, Manitoba (Canada).

**Methods:** There were 169 fecal samples collected between April 18 and June 1, 2018. These included 44 samples of domestic dog feces, 122 of coyote scat, one of fox scat and two of coyote colonic tissue specimens. Samples were frozen (-80°C) for at least 72 hours to inactivate tapeworm ova. Polymerase chain reaction analyses of *E. multilocularis* and *E. canadensis* were performed on all frozen samples.

**Results:** *Echinococcus multilocularis*-positive samples were detected in nine (10.6%) of 85 locations, with one positive sample in a suburban Winnipeg dog park and two positive samples in a popular provincial park. No dog samples were positive for *E. multilocularis*; one sample was positive for *E. canadensis*. In contrast, nine coyote samples (7.3%) were positive for *E. multilocularis* and eight samples (6.5%) were positive for *E. canadensis*. The one fox sample was positive for each. Overall, six samples (3.6%) were positive for both infections.

**Conclusion:** This is the first confirmation of the presence of *E. multilocularis* in coyote feces in the metropolitan area of Winnipeg, Manitoba. In light of the risk this could pose to domestic dogs and human health, periodic surveillance that maps the distribution of this tapeworm could inform the need for additional public health actions.


**Keywords:** alveolar echinococcosis, hydatid disease, zoonotic diseases, coyotes, foxes, *Echinococcus* tapeworms, domestic dogs

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Introduction

Echinococcus species are tapeworms that primarily infect canid species. The two predominant species that have been found in Canada are E. multilocularis and E. canadensis. The sylvatic lifecycle of Echinococcus tapeworms includes coyotes and other canids as definitive hosts and rodents as intermediate hosts. Humans, however, can be incidental (or dead-end) hosts (Figure 1). E. multilocularis can cause a serious disease in humans: alveolar echinococcosis (1,2). This infection, which behaves like a metastatic malignancy, has a high case-fatality rate and requires radical surgical and long-term anthelminthic treatment (3). Although a more benign infection in humans, E. canadensis is also maintained via a sylvatic lifecycle that includes wolves, coyotes, and dogs as definitive hosts and caribou, moose and elk as intermediate hosts. In humans, as incidental hosts, the predominant clinical presentation is cystic hydatid disease primarily in the liver, which may require surgical intervention that carries the risk of severe anaphylaxis should cyst rupture occur (2). These two species are not infrequently found together given they share definitive canid hosts. Echinococcus species tapeworms are asymptomatic and cause no disease in canids, and they are hard to detect, as the worms are only 1–7 mm in length.

How humans become infected

Echinococcus multilocularis ova are quite resistant and can survive and remain infective for nearly one year under favorable conditions (5). As the ova are sticky, they can adhere to the fur of dogs, wild vegetation and garden produce that is grown in feces-contaminated soil (2,6,7). Transmission to humans typically occurs when the hand comes in contact with water, soil, fur or objects contaminated with host feces, followed by inadvertent hand to mouth transfer of the ova. Following human ingestion of E. multilocularis ova, there is a long clinical latency period; from five to 15 years. Early symptoms include abdominal pain, which is followed by jaundice and eventually by severe hepatic dysfunction (2,8). Alveolar echinococcosis is often mistaken for a neoplastic growth, given its propensity for widespread organ infiltration with metastases (2). E. multilocularis may also imitate other diseases, including hepatic carcinoma, cirrhosis and tuberculosis, which may lead to improper diagnostic testing and delayed treatment (3,9,10). Diagnosis is confirmed based on a combination of clinical findings, epidemiological data, imaging, histopathology and/or nucleic acid detection and serology (8). Recommended treatment is radical surgical resection of the parasitic lesion, which in early stages may lead to a complete cure; however, surgical resection can sometimes be incomplete due to the diffuse or undetected spread of the parasite (2,8). As such, post-surgical anthelminthic chemotherapy is recommended for at least two years followed by a minimum of 10 years of close monitoring (8). If untreated, mortality is 90% within 10 years of onset of symptoms (11). Early diagnosis and proper treatment has been shown to improve survival (12).

Echinococcus multilocularis in Canada

Echinococcus tapeworms have been documented in canids in Canada for many years. A study on the prevalence of E. multilocularis in Manitoba canid species, performed 40 years ago in Riding Mountain National Park, found that almost one quarter of all coyotes sampled were infected (13). More recently in 2009, an European strain of E. multilocularis was discovered in central British Colombia in a domestic dog with no history of travel outside Canada, with further research suggesting the possible establishment of this new strain in the local wildlife (14,15). There may be evidence of a similar strain in the wildlife of Saskatchewan (16). Surveys in Calgary and Edmonton published in 2012 determined that E. multilocularis was present in the wild canid populations of those cities (1). Furthermore, in Ontario, recent detections of alveolar echinococcosis in several domestic dogs with no known travel history outside of Ontario suggest that E. multilocularis tapeworms may have established a foci in southwestern Ontario as well (17,18).

Echinococcus tapeworms and cases of human alveolar echinococcosis were first reported in the 1930s in Manitoba and Alberta (19). More recently, in November 2017, a pediatric

Figure 1: Lifecycle of Echinococcus multilocularis

The World Health Organization and the Food and Agriculture Organization of the United Nations have ranked alveolar echinococcosis as the third most important food-borne parasitic disease of global importance (4). As there is no cost-effective way to eliminate E. multilocularis in the sylvatic lifecycle, surveillance is important for human risk assessment.

patient from Manitoba was identified with disseminated alveolar echinococcosis infection (personal communication November 23, 2017, Dr. Sergio Fanella). Similarly, in 2018, several human cases of alveolar echinococcosis were detected in Alberta (personal communication June 5, 2018, Dr. Stan Houston).

With increasing urbanization and sightings of coyotes in urban and suburban areas, *E. multilocularis* may be brought into close proximity to domestic dogs and humans (20,21). In light of these recent human cases, and the sightings of coyotes in and around Winnipeg (20), a survey of wild canids to look for the presence of *Echinococcus* species was conducted.

The aim of this study was to determine the extent of *Echinococcus* species ova, especially *E. multilocularis*, in coyote, fox and domestic dog feces found in the metropolitan area around Winnipeg, and to perform geospatial mapping to determine at-risk areas.

**Methods**

**Collection sites**
The Winnipeg metropolitan area (WMA) was defined as Census Metropolitan Area 602 based on the 2016 census profile from Statistics Canada (22). This consists of the urban area, demarcated by the circumferential perimeter highway surrounding Winnipeg and the peri-urban area that lies outside the perimeter highway and within the WMA. Eighty-five collection sites were identified that covered a wide area of the WMA, including areas frequented by humans and domestic dogs as well as areas with known coyote sightings.

**Sample collection**
Fecal samples were collected between April 18 and June 1, 2018. Samples from domestic dogs were collected differently from the other canid samples. Dog samples were collected from dog parks or submitted directly by dog owners. Two drop-off locations were offered to volunteers who submitted fecal samples from domestic dogs.

Samples from wild canids (mostly coyotes) were predominantly collected by driving along remote roads near coyote habitat based on local knowledge of coyote sightings. Coyote samples were identified via characteristics of the scat sample including shape, texture, presence of hair and bone belonging to small animals as well as supporting features including coyote tracks, nearby coyote dens and previous sightings (20). Similarly the fox scat sample was differentiated by the local conservation officer by its smaller size and the fact there had been known fox sightings nearby.

All fecal samples were picked up individually with plastic bags, sealed and labelled with date of collection, location of collection [Global Positioning System (GPS) coordinate, postal code or address]. For domestic dog samples, the name and phone number of the dog owner was also noted. Two colon samples from carcasses of coyotes were submitted by local wildlife conservation officers who retrieved the colon from two unique animals. Samples were stored at -80°C for at least 72 hours to inactivate *E. multilocularis* ova. The colonic samples were scraped on the mucosal surface to retrieve fecal matter.

**Laboratory testing**
Fecal material was stored and transported in sterile fecal containers, kept on ice, and sent with same day delivery to the reference laboratory (IDEXX Laboratories, Inc.) in Markham, Ontario. Fecal samples were processed immediately upon arrival and total nucleic acid extraction was performed using a KingFisher™ Flex Magnetic Particle Processor (Thermo Fisher Scientific, Vantaa, Finland) with proprietary lysis buffer and magnetic glass particles [Roche Diagnostics, Indianapolis, United States (US)]. Real-time polymerase chain reaction (PCR) assays were carried out using total nucleic acid purified from stool samples on a Roche LightCycler 480™ instrument using the manufacturer recommended cycling protocols and PCR reagents (23). Crossing points were calculated using the second derivative maximum-method analysis module with the high-sensitivity algorithm. Real-time PCR tests included PCR primers and a 6-FAM-TAMRA quenched conventional hydrolysis probe was adapted from Isaksson et al. (24).

The target gene for the real-time PCR tests was the mitochondrial gene *rrnL*. Six quality controls, including PCR positive and negative controls, negative extraction controls, environmental contamination control, spike-in internal positive control and pan-bacterial 16 ssrRNA internal sample control, were run to monitor for extraction efficiency, sample matrix inhibitors and cross contamination in the diagnostic runs. This PCR assay specific for *Echinococcus* species is in the process of being validated and peer reviewed. It was used in this study as it had successfully confirmed a clinical case of *E. multilocularis* of a dog in Ontario (personal communication January 11, 2019, Dr. Roxanne Chan, IDEXX Laboratories, Inc.).

**Analysis**
Data were stored in Microsoft Access and analyzed using Centers for Disease Control (Atlanta, Georgia, US) Epi-Info version 7.2.0.1 for descriptive statistics. Geocoding and mapping were done using Environmental Systems Research Institute ArcMap.

**Results**
In total, 169 samples were collected. This included 122 coyote scat samples, two coyote colon tissue samples, 44 domestic dog fecal samples, of which 34 samples were from unique animals collected directly from dog owners (the other 10 samples were collected from dog parks and may not necessarily be from unique animals), and one fox scat sample.
Echinococcus multilocularis-positive samples were detected in nine (10.6%) of 85 locations, with one positive sample in a suburban Winnipeg dog park and two positive samples in Birds Hill Provincial Park. The other samples appear to be more evenly dispersed across the more rural areas of the WMA. Figure 2 shows the sites where the samples were collected as well as the location of positive samples of E. multilocularis.

Figure 2: Map of the distribution of canid sample collection in the metropolitan area of Winnipeg, indicating Echinococcus multilocularis-positive samples

Of the samples sent for molecular processing, 10 (5.9%) were positive for E. canadensis and 10 (5.9%) were positive for E. multilocularis, with six (3.6%) coinfections in five coyotes and one fox. E. multilocularis was found in 7.3% of all coyote samples, 0% of domestic canine samples and in the single fox sample. E. canadensis was found in 6.5% of all coyote samples, 2.3% of domestic dog samples and in the single fox sample (Table 1).

Table 1: Number and percent of canid samples infected with Echinococcus tapeworms in urban and peri-urban areas of Winnipeg, 2018

<table>
<thead>
<tr>
<th>Source of stool sample</th>
<th>Echinococcus multilocularis</th>
<th>Echinococcus canadensis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td># of samples</td>
<td># positive stools</td>
</tr>
<tr>
<td>Domestic dog(a)</td>
<td>44</td>
<td>0</td>
</tr>
<tr>
<td>Coyote(b)</td>
<td>124</td>
<td>9</td>
</tr>
<tr>
<td>Fox</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Total</td>
<td>169</td>
<td>10</td>
</tr>
</tbody>
</table>

Abbreviation: #, number

\(a\) Of the domestic dog feces samples, 34 were from unique animals, collected directly from dog owners; the other 10 samples were collected from dog parks and may not necessarily be from unique animals

\(b\) The 124 coyote scat samples are not necessarily from unique animals; hence, several samples could have originated from one coyote

Discussion

This is the first confirmation of the presence of E. multilocularis in coyote feces in both urban and peri-urban areas of Winnipeg, Manitoba. The distribution of E. multilocularis in coyotes appears to be wide, with no hotspots. Although E. multilocularis was identified in a popular recreational provincial park and a dog park, no domestic dog samples were positive for this tapeworm. Only one case of the much more benign E. canadensis was found in a domestic dog sample.

Limitations of this study include the short duration of surveillance, possible repeated sampling from the same canids and the use of a molecular assay that is still undergoing peer review (23). No characteristics of the domestic dogs were collected, such as if they were outdoor dogs or consumed rodents. The wild canid scat sampling method was targeted and so does not reflect a uniform collection across the entire region. A much larger sample size and longer surveillance timeframe would be required to more fully map the extent of E. multilocularis canid infection in the Winnipeg metropolitan area.

Implications and next steps

Our findings indicate a risk of human and domestic dog exposure to coyote feces infected with E. multilocularis in the urban and peri-urban areas of Winnipeg.

It is not completely known how often human exposure to the parasite will result in infection leading to alveolar echinococcosis. In 2001, a five-year study in Switzerland was published that showed a high human seropositivity rate to E. multilocularis, with no increase in disease rate. One explanation was a potential increased immunity in this population (25). However, a 2007 review found a significant increase in the incidence of alveolar echinococcosis in Switzerland after 2000, and noted this had been preceded by an increase in this infection in the local fox population 10–15 years prior (26). It was thought this reflected the long clinical latency period of alveolar echinococcosis.
A longer study is now underway to determine if the high prevalence of *E. multilocularis* in the environment is associated with an increased incidence of alveolar echinococcosis in humans (25).

Based on the evidence of *E. multilocularis* in the environment in several provinces and the recent human cases in Alberta and Manitoba, further studies are indicated. To better characterize prevalence and geographic distribution, research with more locations and samples, spatial analysis and in-province molecular diagnostic capabilities are indicated. If warranted, a human seropositivity study of *E. multilocularis* in affected regions could be considered.

This emerging issue is particularly amenable to a One Health approach (27) involving physicians, veterinarians and wildlife experts. Ontario has made *E. multilocularis* infection in domestic dogs and humans reportable, in order to monitor the extent of the infection (28,29). Although currently not notifiable elsewhere, it would be useful to report on the investigation of human alveolar echinococcosis cases, including any travel history (to confirm locally acquired disease versus imported from foreign countries) and information regarding the potential route of exposure to *E. multilocularis* ova. Public education concerning preventative measures to avert domestic dog and human infections may be indicated.

**Conclusion**

*Echinococcus multilocularis* has been documented in Canada for decades in the wild canid population, but human cases have been extremely rare. Over the last two decades, there has been a tendency for wild canids to migrate closer to urban populations, and there have been several recent human cases of alveolar echinococcosis caused by *E. multilocularis*. Although our study did not identify any cross contamination between coyotes and dogs in Winnipeg, the risk of exposure to *E. multilocularis* in urban and peri-urban settings suggests increased awareness and further study would help front line clinicians and public health officials be alert to and monitor this risk.

**Authors’ statement**

CCKT — Project conception, literature searches, study design, field sampling, laboratory investigations, data analysis, writing (original draft, review, and editing)

JB — Project conception, study design, “in kind” laboratory investigations, software support, data analysis, writing (review and editing)

RR — Project conception, study design, software support, spatial data analysis, writing (review and editing)

DD — Project conception, study design, “in kind” laboratory support, data analysis, writing (review and editing)

PJP — Project conception, literature searches, study design, field sampling support, data analysis, writing (review and editing)

**Conflict of interest**

None.

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The results and conclusions are those of the authors and no official endorsement by Manitoba Health, Seniors and Active Living, or Manitoba Agriculture is intended or should be inferred.

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**References**


