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Fatal Cysticercosis by *Taenia crassiceps* (Cyclophyllidea: Taeniidae) in a Presumed Immunosuppressed Canine Host

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ABSTRACT: Cysticercosis in a canine host (*Canis familiaris*) attributable to the taeniid cestode *Taenia crassiceps* is reported for the first time in North America. Numerous parent and daughter cysticerci occurred in a massive intrapleural and intraperitoneal infection in an apparently immunocompromised host. The largest cysticerci were ovoid to elongated, 5–9 mm in maximum length, and armed with 32–34 rostellar hooks in 2 rows; small hooks measured 114–143 \( \mu \text{m} \) long (\( \bar{x} = 124 \pm 8.2 \mu \text{m} \)), and large hooks were 156–180 \( \mu \text{m} \) (\( \bar{x} = 163 \pm 7.4 \mu \text{m} \)). *Taenia crassiceps* is widespread in boreal North America and, like a number of other taeniids, constitutes a potential risk as a zoonotic parasite. The immunological status of the host may be important in determining the outcome of infections for this and other taeniids in atypical hosts.

*A* 6-mo-old female dog weighing 53 lbs was examined for bilateral exophthalmos. The ocular condition was thought to have been caused by an immune-mediated extracocular myositis. Treatment consisting of prednisolone (25 mg p.o., twice a day, PO/BID; Ved Co Inc., St. Joseph, Missouri) and Imuran\textsuperscript{®} (25 mg, PO/BID; e.g., Azathioprine, USP; Glaxo-Wellcome, Research Triangle, North Carolina) for 33 days resulted in an improved condition except for a slight anemia with a packed cell volume (PCV) of 31.9%. Subsequent treatment consisting of prednisolone (20 mg, BID) and Imuran\textsuperscript{®} (25 mg, BID) continued for 5 days; the latter was discontinued when clinical findings included severe ventral abdominal cutaneous bruising, jaundice, and shock with pale mucous membranes and respiratory distress. Clotting time was normal, and PCV was 30%. Serum alkaline phosphatase was 1,963 and the alanine aminotransferase (ALT) was 2,304. Radiographically, the chest and abdomen were unremarkable. A transfusion of 250 ml of canine blood was administered and amoxicillin (500 mg PO/BID; Consolidated Pharmaceuticals, Baltimore, Maryland) was given. Eventually, acute hemolytic crisis and dyspnea developed. Prednisolone (5 mg PO/BID) along with amoxicillin were administered until the time of euthanasia, 6 days later. During this time, PCV dropped from 30 to 17%, the white blood cell count (WBC) from 20,760 to 18,700, ALT from 2,010 to 276, and alkaline phosphatase from 2,548 to 1,806. Presumed immunosuppression in this host is consistent with high dosages of...
prednisolone and Imuran over a 33-day period and the clinical observations of liver failure and changes in the WBC. At the time of euthanasia the dog was icteric, anemic, weak, panting, and anorexic. Euthanasia was accomplished by administration of Socumb® (pentobarbital sodium at 6 gm/ml; The Butler Company, Columbus, Ohio).

During postmortem, numerous (too abundant to count accurately), ovoid to slightly elongate cysticerci, approximately 5–9 mm in length for the largest metacestodes, were found free and observed moving within the pleural and peritoneal cavities (Figs. 1, 2). The lungs were diffusely edematous and with multiple hemorrhages. Additionally, multiple cavities, each con-
taining 1 cysticercus or multiple cysticerci, were scattered throughout the lungs. Other tissues or organs were not involved, and otherwise appeared unremarkable. Strobilate specimens of cestodes were not present in the small intestine.

Samples of various tissues, including extraocular muscles, and numerous cysticerci were collected and fixed in buffered 10% formalin. Fixed tissues were trimmed, processed, embedded in paraffin, and sectioned. Tissue sections, 4–5 μm thick, were stained with hematoxylin and eosin (H&E). Representative metacestodes were stained with Semichon’s acetic carmine, dehydrated in ethanol, cleared in xylene, and mounted entire in Canada balsam; hooks from some specimens were mounted separately to allow detailed study of structure.

Histologically, extraocular muscles had extensive infiltration of lymphocytes and plasma cells between myofibers and multifocal necrosis and loss of myofibers. Proteinaceous fluid, blood, and many macrophages were present in alveoli. Metacestodes were present within capsules delimited by fibrous connective tissue containing eosinophils, lymphocytes, and plasma cells (Figs. 2, 3).

Cysticerci were morphologically and ontogenetically consistent with *T. crassiceps*. Among 10 specimens collected from the peritoneal cavity and mounted entire, all were ovoid to elongate and 5–9 mm in maximum length by 3–4 mm in width; numerous specimens were considerably smaller. Cysticerci exhibited exogenous and endogenous budding at the polar end opposite the invaginated scolex (Fig. 4); development was proliferative. Among 5 scolices, the rostellum was armed with 2 rows of 16–17 small and large hooks (Figs. 5, 6), respectively, for a total of 32–34. Small hooks measured 114–143 μm long (n = 35; $\bar{x} = 124 \pm 8.2$ μm), and large hooks were 156–180 μm (n = 35; $\bar{x} = 163 \pm 7.4$ μm). Representative specimens have been deposited in the U.S. National Parasite Collection, Biosystematics and National Parasite Collection Unit, USDA, Agricultural Research Service, Beltsville, Maryland (USNPC 87835).
**Taenia crassiceps** is unique among the taeniids in having a proliferative cysticercus that develops asexually by exogenous budding (Freeman, 1962). Rostellar hooks have a characteristic form, with the blade being markedly longer than the handle (Abuladze, 1965; Verster, 1969). In strobilate specimens in canids and in mature metacestodes in rodents, there are usually 30–36 hooks (occasionally as few as 28), with small hooks measuring 107–155 μm and large hooks 155–200 μm in length (Rausch, 1959; Freeman, 1962; Leiby and Whittaker, 1966; Verster, 1969). Chermette et al. (1996) reported 31–34 hooks, measuring 92–153 μm and 145–201 μm for small and large forms, respectively, from cysticerci infecting dogs in France.

Globally, cysticercosis or coenurosis due to species of *Taenia* Linnaeus, 1758 in companion animals has been reported sporadically, and most infections have been documented in feline hosts (e.g., Kingston et al., 1984; Huss et al., 1994). Reports of cysticercosis other than *Taenia solium* Linnaeus, 1758 in canine hosts are apparently rare, although dogs serve as definitive hosts for at least 7 species of *Taenia* (Verster, 1969; Chermette et al., 1993, 1996). In the United States, only cysticerci of *T. pisiformis* (Bloch, 1780) have been reported in dogs (Ivens et al., 1969); the record herein is the first from North America for cysticercosis attributable to *T. crassiceps* in a canine host. In France there have been 3 reported cases of cysticercosis due to *T. crassiceps* in a canine host. Among these hosts, 2 had cysticerci in subcutaneous tissues and 1 had a massive peritoneal infection similar to that observed in the current study.

The epizootiology of canine cysticercosis is incompletely understood. All cases have been in rural situations where foxes and a diversity of rodents are common (Chermette et al., 1996). In Michigan and other boreal regions of the U.S., definitive hosts for *T. crassiceps* would be expected to include red fox, coyote, and dogs, whereas intermediate hosts may be represented by voles and mice (e.g., species of *Microtus* Schrank and *Peromyscus* Gloger), squirrels (e.g., species of *Tamias* Illiger, *Sciurus* Linnaeus, and *Tamiasciurus* Trouessart), woodchucks (e.g., *Marmota monax* (Linnaeus)), and other rodents (see Freeman, 1962). With respect to the current case, the locality is an isolated area of northern Michigan, red foxes and coyotes are known in the local area, and the dog had recently eaten mice that had been caught by resident cats. Other companion animals in the household included 2 domesticated cats that were usually outside the residence.

Exposure to infection for canine hosts may follow several potential routes: (1) via eggs in the environment, (2) autoinfection via eggs from a strobilate and gravid adult, and (3) via ingestion of cysticerci in the intermediate host (Freeman, 1962; Kroze and Freeman, 1982; Chermette et al., 1993, 1996). Relatively few oncospheres may be expected to result in a massive infection (Freeman, 1962), and there may be up to 13,000 eggs in a typical gravid segment (Miyaji et al., 1990). In the rodent intermediate host, complete development of the scolex requires 35–42 days; budding occurs by 28 days PI, leading to massive infections that result from the cumulative production of cysticerci (Freeman, 1962).

The presence of numerous parent and daughter cysticerci suggests that the dog was infected for at least several months or had initially received a large infective dose. Alternatively, the stage of development for metacestodes and the occurrence of both parent and daughter cysticerci in a young dog may be consistent with an infection acquired following ingestion of an intermediate host.

Previous disease, stress, or medical and surgical procedures leading to an impaired immune system may predispose canines to cysticercosis (Chermette et al., 1993, 1996). Thus, presumed immunosuppression of the dog in the current study, due to administration of prednisolone and Imuran may have been a contributing factor to a massive proliferative infection by *T. crassiceps*; Imuran is typically used as an immunosuppressant (Coppoc, 1988). Immunosuppression of this host is clearly suggested based on continuous treatment for 33 days at the highest recommended dosages for these drugs. The immunological status of the host may be important in determining the outcome of infections for this and other taeniids.

*Taenia crassiceps* is widespread in boreal North America and, like a number of other taeniids, constitutes a potential risk as a zoonotic parasite (Freeman et al., 1973). Infections by *T. crassiceps*, however, may be particularly serious due to their proliferative nature, in contrast to cysticercosis associated with other species of *Taenia*. Subcutaneous infections may pose a limited clinical problem in humans, but localization in other organs could result in serious disease (Freeman et al., 1973; Chermette et al., 1996). The recent reports from Europe of proliferative infection by *T. crassiceps* in immunocompromised patients with HIV highlights the potential importance of taeniids as zoonotic parasites (Chermette et al., 1995).

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**LITERATURE CITED**


