



Revista Brasileira de Parasitologia
Veterinária

ISSN: 0103-846X

zacariascbpv@fcav.unesp.br

Colégio Brasileiro de Parasitologia
Veterinária
Brasil

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Revista Brasileira de Parasitologia Veterinária, vol. 26, núm. 1, enero-marzo, 2017, pp. 47
-53

Colégio Brasileiro de Parasitologia Veterinária
Jaboticabal, Brasil

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Different inoculum loads of *Toxoplasma gondii* induce reduction of myenteric neurons of the rat colon

Diferentes inóculos de *Toxoplasma gondii* induzem a redução de neurônios mioentéricos no cólon de ratos

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Received October 14, 2016

Accepted January 17, 2017

Abstract

Toxoplasmosis, a disease caused by *Toxoplasma gondii*, is an important health problem, especially in immunocompromised hosts. *T. gondii* uses the gut wall as an infection gateway, with tropism for muscular and nervous tissues causing intestinal alterations, including some in the enteric nervous system. This study aims at investigating the colon of rats infected by *T. gondii* in order to understand how the amount of oocysts influences in myenteric neuronal changes. Sixty Wistar rats (*Rattus norvegicus*) were divided into six groups. One group remained as a control and the others received inocula of 10, 50, 100, 500 or 5,000 oocysts of *T. gondii*. The animals were euthanized after 30 days of infection. The total neuronal population and the nitrergic subpopulation in the colon myenteric plexus of each animal was counted. The data were statistically analyzed showing less weight gain in rats with 10, 500 and 5,000 oocysts. A decrease in the number of total neurons with 50, 100 or 5,000 oocysts and an increase in the nitrergic population with 10, 100, 500 or 5,000 oocysts were verified. These results show that neuronal alterations are more significant when the infection is induced by larger inocula and reinforces the suspicion that neuronal loss is directed at cholinergic neurons.

Keywords: Enteric nervous system, myenteric plexus, ascending colon, toxoplasmosis.

Resumo

A toxoplasmose, doença causada pelo *Toxoplasma gondii*, é um importante problema de saúde, principalmente em imunocomprometidos. *T. gondii* utiliza a parede do intestino como porta de entrada no hospedeiro e tem tropismo pelos tecidos muscular e nervoso provocando alterações intestinais, inclusive no sistema nervoso entérico. Este estudo buscou analisar o cólon de ratos infectados por *T. gondii* para entender como a quantidade de oocistos influencia nas alterações neuronais mioentéricas. Foram utilizados 60 ratos Wistar (*Rattus norvegicus*) em seis grupos. Um dos grupos permaneceu como controle e os demais receberam inóculos de 10, 50, 100, 500 ou 5.000 oocistos de *T. gondii*. Os animais foram submetidos a eutanásia após 30 dias de infecção. No plexo mioentérico do cólon dos animais foram quantificadas a população neuronal total e a subpopulação nitrérgica. Os dados foram analisados estatisticamente demonstrando inferior ganho de peso nos ratos com 10, 500 e 5.000 oocistos. Verificamos diminuição no número de neurônios totais com inóculos de 50, 100 ou 5.000 oocistos e aumento da população nitrérgica com 10, 100, 500 ou 5000 oocistos. Estes resultados mostram que alterações neuronais são mais significativas quando a infecção é induzida por inóculos maiores e reforça a suspeita de perda neuronal direcionada a neurônios colinérgicos.

Palavras-chave: Sistema nervoso entérico, plexo mioentérico, cólon ascendente, toxoplasmose.

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Introduction

Toxoplasma gondii is an obligate intracellular coccidian protozoan parasite, which is widely distributed (WEISS & KIM, 2007). In Brazil, seroepidemiological studies show the presence of antibodies against *T. gondii* in about 65% of the population (FEREZIN et al., 2013; REICHE et al., 2000; SARTORI et al., 2011). The definitive hosts in the life cycle of *T. gondii* are cats and other felids, and several animal species such as birds and mammals act as intermediate hosts (DUBEY, 2007). In humans, its main transmission probably occurs through the ingestion of water or food contaminated with oocysts (MONTROYA & LIESENFELD, 2004; WEISS & KIM, 2007). After this, the oocysts release the sporozoites, upon crossing the intestinal wall, change into their infective form, tachyzoites, which reach the bloodstream (DUBEY, 2007).

The pathological process of toxoplasmosis depends on the parasite genotype. One of the major and classical genetic classifications of *T. gondii* strains in Europe and USA comes from the SAG2 gene analysis (HOWE & SIBLEY, 1995). The type I strain is considered of high virulence, type II is of intermediate virulence and type III is of low virulence (SILVA et al., 2005; FREYRE et al., 2001). The strain used in this study, identified as ME49 (genotype II), was isolated, in the northern hemisphere, from the muscles of a ram (LUNDE & JACOBS, 1983), and the fact that it is not of high virulence favors its use in studies of the chronic phase of the infection (MAUBON et al., 2008).

In the chronic phase of the disease, the protozoan tends to form tissue cyst tropism in the nervous and muscle tissues. The intestinal wall is the first barrier it transposes to infect the host and then to lodge in these sites (DUBEY et al., 1998). Previous studies of our research group focused on the changes caused by *Toxoplasma gondii* infection in the enteric nervous system (ENS), which is responsible for controlling important functions of the gastrointestinal tract (GIT) (BONAPAZ et al., 2010; ODORIZZI et al., 2010; PEREIRA et al., 2010; HERMES-ULIANA et al., 2011; ARAÚJO et al., 2015; GÓIS et al., 2016a; TREVIZAN et al., 2016).

The ENS consists of enteroglia cells and neurons distributed in two ganglionic plexuses: the myenteric plexus, which is located between the longitudinal and circular muscle layers, and mainly controls the contraction and the intestinal peristalsis; and the submucosal plexus, situated along the submucosa, responsible mainly for the regulation of mucus secretion and local blood flow (FURNESS, 2006; GÓIS et al., 2016b). Among the different neuronal populations within ENS, the nitrergic population comprises neurons with nitric oxide synthase type I activity (NOS-I), a NADPH-dependent enzyme involved in the synthesis of nitric oxide. Among its numerous functions, nitric oxide acts as a nerve cell transmitter, promoting the relaxation and inhibiting the contraction of the intestinal smooth muscle (DUSSE et al., 2003; IYENGAR et al., 1987).

Recent reports indicate that rats infected with *T. gondii* present histological and immunological changes in the GI similar to those seen in inflammatory bowel diseases (LIESENFELD, 2002; SCHREINER & LIESENFELD, 2009). Thus, the study of changes caused by *T. gondii* infection in the ENS may help shed

light on the changes that occur in those pathological conditions. Although the colon is the main organ involved in inflammatory bowel diseases (MEYERS & JANOWITZ, 1989; BAUMGART & SANDBORN, 2012), and is associated with the cause of intestinal tract disorders in various diseases, only a few studies have examined the relationship between *T. gondii* infection and local innervation. Sugauara et al. (2008) found that rats infected with a Brazilian strain of *T. gondii*, BTU-II (genotype III), for 30 days exhibited myenteric neuron hypertrophy of the colon. Another study, which involved genotype I tachyzoites (BTU-IV strain), found an increase of cytoplasm in this neuronal population (SOARES et al., 2009). However, neither of the two aforementioned studies reported quantitative changes, and there are no reports of studies with the ME49 strain or any other of genotype II.

Therefore, the purpose of this study was to evaluate the changes caused by different inoculum loads of *T. gondii* on total myenteric and nitrergic neurons in rat colon, and to help consolidate a standard experimental model of infection with *T. gondii* that best represents its effects on the GIT.

Materials and Methods

The experimental protocol was approved by the Ethics Committee for Animal Experimentation of the State University of Maringá (Protocol No. 081/2012).

Experimental design

Sixty 60-day-old Wistar rats (*Rattus norvegicus*) were randomly distributed into six groups of 10 rats each. The control group (CG) received sterile saline orally by gavage, while the animals of the other groups were ranked according to the number of inoculated *T. gondii* oocysts (10, 50, 100, 500, or 5000). Inoculation was also performed orally, using the ME49 strain (genotype II) oocysts, previously sporulated and resuspended in 1 mL of sterile saline, which was obtained from the Department of Preventive Veterinary Medicine, State University of Londrina, Paraná, Brazil.

The animals were housed in individual cages for 30 days, with 12-hour light/dark cycles, under controlled temperature and humidity, and were fed rodent chow and water *ad libitum*. The animals were weighed on days 1 and 30 to calculate their body mass, and were observed during the experimental period for possible clinical signs of toxoplasmosis infection. The experimental design was entirely randomized.

Confirmation of infection with *T. gondii*

Thirty days after inoculation, blood samples were drawn from all the animals and subjected to a direct agglutination test to detect IgG antibodies against *T. gondii*, indicating immunity to the protozoan. Samples with a titer $\geq 1:25$ were considered positive (DESMONTS & REMINGTON, 1980).

Euthanasia and sample collection

Thirty days after inoculation, the animals were euthanized by deep anesthesia with halothane vapor (VIVAS et al., 2007). Each animal was laparotomized, followed by the removal and measurement of the ascending colon to calculate the area, and the removal of two segments of approximately 3 cm for subsequent histological processing.

Sample processing

The myenteric plexus was marked using two techniques: the pan-neuronal Giemsa staining technique (BARBOSA, 1978) and the NADPH-diaphorase (NADPH-d) histoenzymological technique, which marks the neuronal subpopulation with NOS-I activity (PHILLIPS et al., 2003; SCHERER-SINGLER et al., 1983).

Giemsa technique

The segments were washed with a 0.9% NaCl solution and filled with formaldehyde-acetic fixative solution for at least 48 hours. The mucosa and submucosa were then removed using a stereomicroscope with trans-illumination to reveal the myenteric plexus, which was then stained with methylene blue in Sorensen's phosphate buffer solution (pH = 7.0) for 24 hours (BARBOSA, 1978). The whole mounts were then dehydrated in a series of alcohol solutions of increasing concentration and cleared in xylol.

NADPH-diaphorase technique

The segments were washed and filled with 0.1M PBS (pH 7.3), fixed with 4.0% paraformaldehyde in 0.1M phosphate buffer for 30 minutes, and immersed in PBS with 0.3% Triton X-100 for 10 minutes. They were then washed in ten times in 10-min baths in PBS for subsequent incubation for two hours in a medium containing the following ingredients in 200 mL: 100 mg of β -NADPH (Sigma, St. Louis, MO, USA), 0.6 mL of Triton X-100, and 50 mg of NBT in Tris-HCl solution (Life Technologies, Grand Island, NY) (SCHERER-SINGLER et al., 1983). The membrane preparations were then dehydrated in series of alcohol solutions of increasing concentration and cleared in xylol.

Quantitative analysis of myenteric neurons

All the myenteric neurons visible under 400X magnification in 120 fields of an optical microscope (Motic) were counted. The number of neurons of intestinal area was calculated based on the measurement of the microscopic field (0.249 mm²). The same procedure was performed for both preparation techniques in all the animals.

Statistical analysis

The sample residues were submitted to the Shapiro-Wilk test to verify distribution type (ZAR, 1999). Data with normal distribution were expressed by mean \pm SDs. Results of the neuron counts

analysis were compared between groups using one-way analysis of variance (ANOVA) and Dunnett's multiple comparison post hoc test. The statistical analysis of the data was performed using Graph Pad Prism 5.01 software. For all statistic tests, p-values less than 0.05 were considered significant.

Results

The animals of the CG showed negative IgG serology against *T. gondii*, while the animals inoculated with the parasite presented positive serology, albeit without obvious clinical signs of toxoplasmosis, such as changes in the intestinal tract. The animals of the infected groups, G10, G500 and G5000, presented lower body mass gain than the CG (p<0.05), although the colon area in all the groups showed no difference (p> 0.05).

The neuron counts revealed a significant decline in the total neuronal population of groups G50, G100 and G5000 (Figure 1A), while the nitrergic subpopulation (NADPH-diaphorase positive or NADPH-d⁺) showed an increase in G10, G100, G500 and G5000 (p<0.05) (Figure 1B).

The Giemsa technique (Figure 2A-C) revealed a decline in the number of neurons in the myenteric plexus in the colon of rats infected with 50, 100 or 5000 oocysts. A comparison of the number of myenteric neurons of each animal in the infected groups and the average number presented by the CG and its confidence interval by both techniques indicated that 83% of the animals in G100 and 86% in G5000 presented neuronal loss. These data are consistent with the statistical results, which showed a significant decrease in the average number of neurons. However, all the infected groups had animals that presented this result, particularly G50, in which 67% of the animals showed decreasing numbers. Unlike what was observed in the total neuronal population, the NADPH-d⁺ technique (Figure 2D-F) revealed a quantitative increase in the nitrergic subpopulation. An individual analysis indicated a tendency for the number of NADPH-d⁺ neurons to increase, particularly in 86% of the animals in G500 and 80% of those in G100. Figure 3 depicts individual variations in the animals in each group that presented an increase or decrease in the number of neurons.

Discussion

Despite the established infection, changes in the population of enteric neurons were observed mainly in larger inoculum loads, and reveals the total population of neurons and that the average area of the colon in the infected groups remained unchanged, neuronal loss related to *T. gondii* infection probably occurred.. This result may be due to the action of chemical mediators that are associated with the inflammatory response triggered by *T. gondii*, including nitric oxide (NO) and IFN- γ , which participate in the mechanism of apoptotic induction and are linked to morphological and quantitative changes of enteric neurons in protozoal infections (ARANTES et al., 2004; NISHIKAWA et al., 2007). On the other hand, other studies of the colon of rats infected with *T. gondii* found no changes in the total number of neurons of the myenteric plexus (SOARES et al., 2009; SUGAUARA et al.,

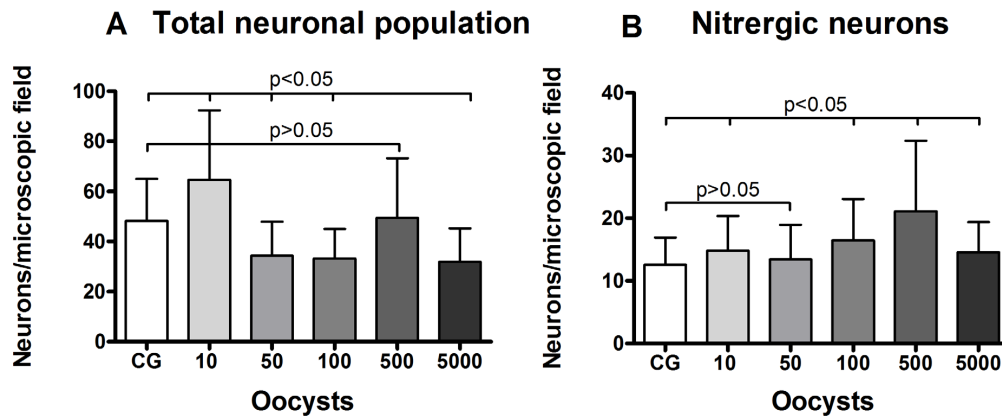


Figure 1. Number of myenteric neurons of the colon of rats infected with different *Toxoplasma gondii* inoculum loads, expressed in neurons/microscopic field by the Giemsa (A) and NADPH-diaphorase techniques (B). ANOVA followed by Dunnett's multiple comparison post hoc test.

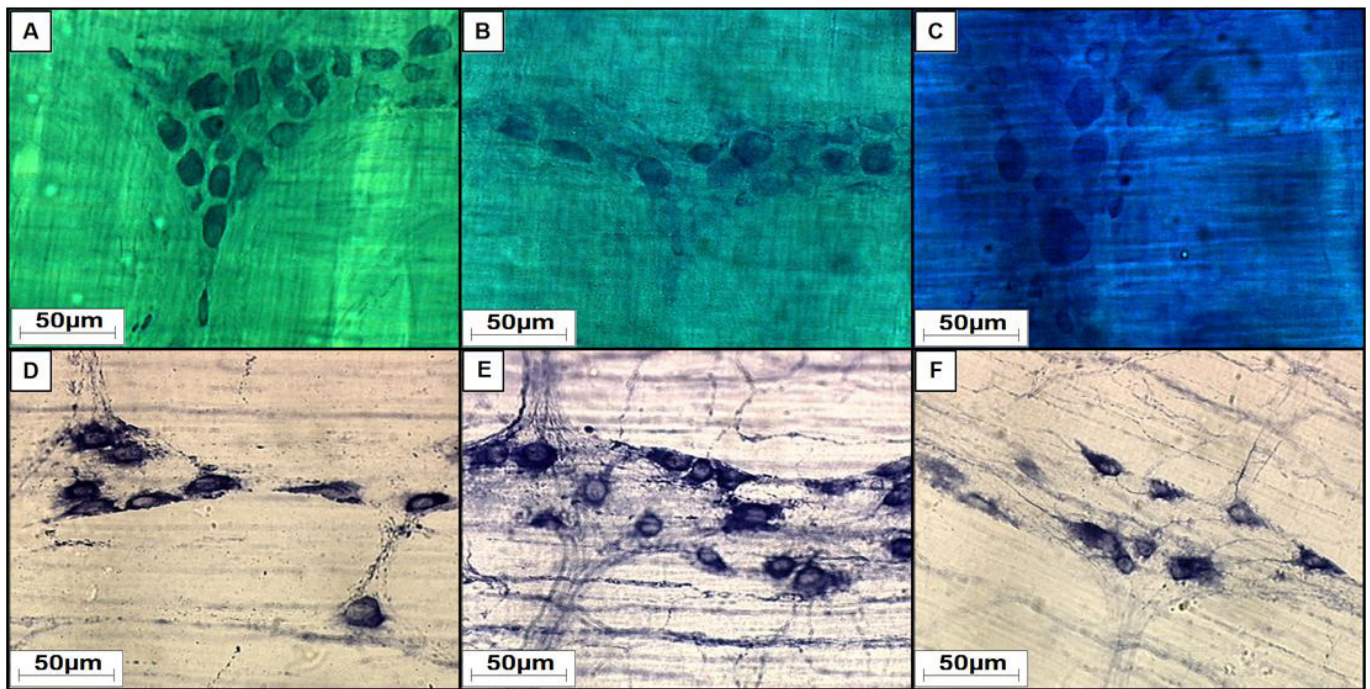


Figure 2. Photomicrographs showing different neuronal densities by the Giemsa (A=CG, B=G100 and C=G5000) and NADPH-diaphorase (D=CG, E=G500 and F=G5000) techniques. 400X.

2008). This divergence in findings is explained by differences in the experimental protocol, such as the evolutionary form of the inoculated parasite and the genotype of the strain. A study of the small intestine in which a similar experimental protocol was used revealed neuronal loss with inoculum loads of 10, 100, 500 and 5000 oocysts, demonstrating a greater depletion effect of the infection on neurons in the initial part of the intestines (VICENTINO-VIEIRA et al., 2015). This can be explained by the natural presence of a larger number of intraepithelial lymphocytes in the small intestine, resulting in a more intense immune

response with greater release of cytokines and other inflammatory mediators, and hence, greater tissue damage (BEAGLEY et al., 1995). The release of *T. gondii* sporozoites also reportedly occurs in the small intestine (FERGUSON & DUBREMETZ, 2007), which appears to trigger a more intense inflammatory response in this portion of the GIT, also suggesting that neuronal loss seems not to be caused by direct action of the parasite.

Due to the complexity of the biological systems, individuals within the same group respond differently, despite all the care taken to control the variables. It should be noted that the neuronal

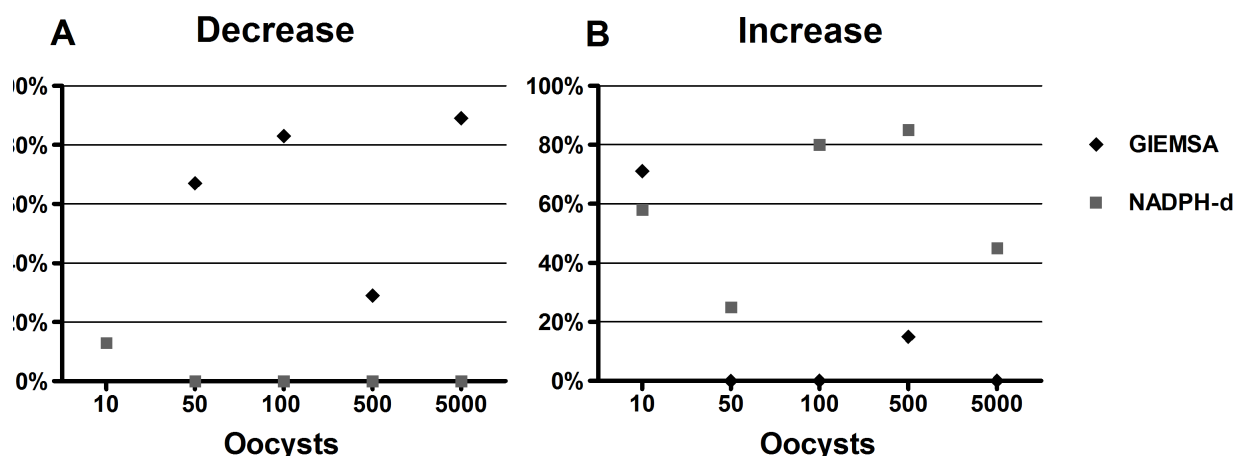


Figure 3. Percentage of animals in groups 10, 50, 100, 500 and 5000 oocysts that showed a decrease (A) and an increase (B) in neuronal counts compared to Control group, by the Giemsa and NADPH-diaphorase techniques.

assessment involved 120 different and random microscopic fields and that the quantitative analysis was blind.

These analyses reinforce the belief that nitrergic neurons are more resistant to apoptosis (COWEN et al., 2000; PHILLIPS et al., 2003), and that after contact with *T. gondii*, a portion of the total surviving neuronal population begins to express the enzyme NADPH-diaphorase, and therefore NOS-I, in addition to those they already expressed under normal conditions. This phenomenon may indicate a relationship between the immune response against the parasite and the production of nitric oxide in the myenteric plexus. This is corroborated by similar studies of the jejunum of these animals (VICENTINO-VIEIRA et al., 2015) and that of pigs (ODORIZZI et al., 2010), in which a genotype III strain was used. However, no studies were found about the effects of toxoplasmosis infection on the NADPH-d⁺ neurons specific to the colon.

The muscle layers of the intestine are densely innervated by the nerve fibers of the myenteric plexus. The neurons in this plexus can be divided into two different functional types: inhibitory, whose transmission, called non-adrenergic non-cholinergic (NANC), is identified by the concomitant presence of the enzyme NOS and vasoactive intestinal peptide (VIP), and excitatory motor neurons, which are revealed by immunoreactivity to tachykinins and to the vesicular acetylcholine transporter, which are considered cholinergic neurons (DE GIORGIO et al., 1994; FERRI et al., 1983; LLEWELLYN SMITH et al., 1984). Inhibitory and excitatory neurons innervate the muscle with different fibers (WATTCHOW et al., 1988).

The increase in the NOS⁺ neuron population, occurring simultaneously with the reduction of the total population, leads us to suppose that the myenteric neuronal depletion caused by toxoplasmosis infection was concentrated in the cholinergic neurons, causing a downward trend in the contractility of the fibers of the intestinal smooth muscle, a phenomenon that may lead to motility disorders. Less contractile muscle fibers lead to slower intestinal flow and constipation, although this sign was not

detected in the rats of the infected groups. Because the ME49 strain is not highly virulent, the tissue damage and possible neuronal loss may have been insufficient to trigger noticeable changes during the infection period under study.

In this study, we concluded that the depletion effect of the genotype II strain of *T. gondii* on the myenteric neurons of the rat colon is significantly achieved through oral inoculation with 5,000 oocysts, and that this effect seems to occur mainly on the cholinergic population of these cells. An increase in the NADPH-d⁺ subpopulation was also found in response to the inoculation of 10, 100, 500 or 5,000 oocysts. This experimental model proved to be suitable for studies of the chronic phase of *T. gondii* with an infection time of 30 days.

To help elucidate the phenomena found in this work, additional studies are needed to assess changes in the intestinal tract. Further research is also needed to assess not only quantitative effects but also atrophy and hypertrophy of the myenteric neurons of the colon in infections with larger numbers of oocysts, employing techniques that reveal both nitrergic and cholinergic subpopulations. Studies aimed at clarifying the role of cellular immune response and its mediators, i.e., intraepithelial lymphocytes, macrophages, IFN- γ , NO, cytokines, etc., are already ongoing and may help shed light on the effects of *T. gondii* infection in the myenteric plexus and in the other structures of the intestinal wall.

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