Neurocysticercosis (NCC) is one of the main parasitic diseases of the central nervous system (CNS) caused by Taenia solium cysticerci, which are responsible for significant morbidity associated with seizures and hydrocephaly.1-3 It is considered an emerging disease in most developed countries especially due to the increase in the number of immigrants from endemic areas and to a few documented cases of local transmission.3,4 According to a recent study, with patients from a neurologic clinic in Texas, USA, 53% of the hospitalizations occurred due to NCC, and epilepsy was the main clinical symptom. From these NCC patients, 93% were Hispanic immigrants.4 As it presents great clinic diversity, NCC has been the target of several clinic, serum epidemiologic and histopathologic studies, which aim at understanding the mechanisms involved in the immune and inflammatory responses.6

The use of experimental models to the comprehension of the host-parasite relationship has become an excellent tool to...
study the human cysticercosis in several organs, including its severe form, the NCC. The parasite most used in those cysticercosis experimental models is Taenia crassiceps as it presents rapid developing cycle, easy maintenance, and antigenic similarity to T. solium. The intraperitoneal model is the most diffused one as it is useful on the evaluation of genetic factors involved in the resistance or susceptibility of hosts and especially of the immunological mechanisms. The most described experimental model in literature to NCC studies is the one reported by Cardona et al., who used Mesocestoides corti.

According to the literature, BALB/c mice are less resistant than C57BL/6 to the experimental intraperitoneal infection with T. crassiceps cysticerci. While in the most severe form of the disease, NCC, there are few reports on the inflammatory and immunological mechanisms from the host as well as on the evasion mechanisms used by the parasite as to survive in this hostile environment. It is known that, in the beginning of the intraperitoneal infection, there is a type 1 cytokine that cause an inflammatory response, which controls the parasite’s growth. However, this response rapidly changes to type 2 or even to a mixed type 1/type 2 profile of cytokines, which is permissive to the parasite’s growth. The type 2 immune response results in a derestrict growth of the parasite that may lead to the death of the animal in experimental cases, demonstrating little or no immunologic resistance to the parasitary growth.

The objectives of this study were to develop an experimental model that could induce NCC in mice by using T. crassiceps cysticerci, to observe the differences in the inflammatory reaction and consequent lesions in BALB/c and C57BL/6 mice, and to determine their susceptibility or resistance to the infection.

**METHODS**

**Maintenance of the parasite**

The biological cycle of T. crassiceps (ORF strain) has been maintained in the animal facility of the Tropical Pathology and Public Health Institute from the Federal University of Goiás (IPTSP/UFG), since 2002. Ten initial phase cysticerci were inoculated in the intraperitoneal cavity from 8 to 12 weeks-old female BALB/c mice, where they were multiplied by budding. Approximately 90 days after inoculation, the animals were euthanized and necropsied, and the initial stage cysticerci were removed and washed several times with sterile saline solution and inoculated in the peritoneal cavity of noninfected BALB/c female mice.

**Animals**

Matrices from conventional female BALB/c and C57BL/6 mice were maintained in the animal facility of the IPTSP/UFG. For this study, we used animals from 8 to 12 weeks-old and with 20 to 30 g. These animals were intracranially inoculated with three to five initial stage cysticerci. The animals were divided into four groups, containing five animals for each experimental day (7, 30, 60, and 90 days after the inoculation) named as: Group 1 – BALB/c mice infected with cysticerci; Group 2 – BALB/c mice inoculated with sterile saline solution; Group 3 – C57BL/6 mice infected with cysticerci; and Group 4 – C57BL/6 mice inoculated with sterile saline solution.

The ethical principles for animal experimentation professed by the Brazilian Society of Laboratory Animal Sciences (SBCAL) were followed. This study was authorized by the Ethics Research Committee of the Federal University of Goiás (CoEp/UFG), registration number 034/09.

**Experimental infection**

The animals were weighted and anesthetized previously to the inoculation. The anesthetics consisted of a solution of Ketamine (100 mg/mL) and Xilazine (20 mg/mL) in the proportion of 0.1 mL/10 g. After the trichotomy of the head superior portion and the antisepsis with topic iodine, a longitudinal and median incision was made on the skin of the skull with a scalpel. The trepanation orifice was performed with a drill (4.5x2 mm) moved by a micromotor (LB100-Belttec) in the topography of the right parietal bone at 3 mm, from the median line (sagittal suture), and at 3 mm posterior to the coronal suture and with 4 mm of depth. The infected animals were intracranially inoculated with cysticerci and afterwards the trepanation orifice was closed with sterile dental alginate and the incision was sutured.

**Removal of the encephala**

At 7, 30, 60, and 90 days after the inoculation (DAI), the animals were intraperitoneally anesthetized with 0.1 mL/10 g of the Xilazine 2% and Ketamine 10% solution. Afterwards, the animals were euthanized by cervical dislocation and had their encephala removed to posterior analysis.

**Histopathological analysis**

The histopathological analysis was performed with fragments of the encephala. They were fixed with 3.4% buffered formalin; dehydrated with alcohols; clarified with Xylol; and then blocked into paraffin and sectioned into 5 micrometer sections; stained with hematoxylin and eosin (HE) and other histochemical techniques when necessary, such as picrosirius, to fibrosis identification, periodic acid of Schiff (PAS), to glycric radicals identification, von Kossa, to calcium salts deposits identification, and Congo red to amyloidosis identification.

**Quantification of the cysticerci and their classification into development stages**

Through microscopic analysis, the anatomical localization of the cysticerci was described as well as its classification, according to its development stage into initial, larval, and final.
General pathologic processes analysis

The general pathologic processes were analyzed inside the parasite, in the host-parasite interface, and in the host tissue considering the presence of an inflammatory process (parenchyma, meninges, and blood vessels) and its cellular profile, edema, hyperemia, calcification, fibrosis, glycidic radical deposits, gliosis, and microgliosis. The pathologic processes described were classified into a semi-quantitative way according to the following criteria: absent; discrete with up to 25% of the compromised area; moderate from 26 to 50% of the compromised area and accentuated above 50% of the compromised area

Statistical analysis

The statistical analysis was performed by using the Sigma Stat 3.5 program. All variables were tested as to their normal distribution and homogenous variation. As they presented non-normal distribution, the variables were analyzed by the nonparametric Mann-Whitney’s test. The differences were considered significant when p<0.05.

RESULTS

This study described the development of an experimental model to NCC studies involving two mice lineages. These lineages were compared as to their susceptibility or resistance to infection by T. crassiceps cysticerci, and also as to the inflammatory process that each one developed throughout 90 days of intracranial infection.

All infected animals presented cysticerci located in the brain ventricles (lateral and dorsal) provoking inflammation, expansion to this cavity and, consequently, deviation of the median line. Those effects were observed with greater intensity in BALB/c mice (Fig 1 and 2, Tables 1 and 2).

On both lineages, a decrease in the consistency and hypotrophy of the encephala parenchyma was observed. At 7 DAI, the cysticerci found were classified into the initial development stage on both lineages, while at 30 DAI larval stage cysticerci were found only in C57BL/6 mice (Fig 3A to D). Final stage cysticerci were found more precociously in C57BL/6 mice, at 60 DAI (Fig 3E and F) than in BALB/c mice, at 90 DAI (Fig 3G, Table 1). Also, the inflammatory infiltration observed in BALB/c mice being more intense than the one observed in C57BL/6 mice (Fig 3H, Table 1). Final stage cysticerci also presented discrete areas of fibrosis, decreased glycidic material deposit, and necrosis.

When analyzing the host-parasite interface from infected BALB/c mice, it was possible to observe an intra-parenchymatous inflammatory infiltration with predominance of polymorphonuclear cells. In C57BL/6 mice, the inflammatory cells predominance was of mononuclear cells. At 30 DAI the perivasculitis was significantly greater in BALB/c mice.
Microgliosis was observed at 7 DAI in C57BL/6 mice, while in BALB/c mice it was observed only at 30 DAI. Also, in the first ones a more intense gliosis in relation to the latter was observed. Discrete areas of calcification were found in the brain parenchyma in the region of the caudal putamen in infected C57BL/6 mice at 90 DAI (Table 3).

**DISCUSSION**

Both lineages of mice used in this study could induce necrosis on the parasites at 90 DAI. However, in C57BL/6 mice, this destruction occurred in a more efficient manner since 60 DAI. In spite of that, the inflammation composed mainly by
polymorphonuclear cells associated with edema, perivascularitis, and meningitis in BALB/c mice was significantly greater than what was observed in the C57BL/6 ones during the experimental period. Fragoso et al.\textsuperscript{23}, when inoculating BALB/c mice via intraperitoneal with \textit{T. crassiceps} cysticerci, observed that they presented a greater parasitary burden than C57BL/6 mice and considered the first susceptible to \textit{T. crassiceps} cysticerci infection. We believe that this lower capability to induce necrosis on the parasites from BALB/c mice may have occurred due to the predominance of polymorphonuclear cells in the inflammatory infiltration throughout the infection. On the other hand, C57BL/6 mice already presented mononuclear cells in the inflammatory infiltration at 60 DAI. Probably due to this difference, the intensity in the ventriculomegaly, the destruction of the adjacent parenchyma, the deviation of the median line, and the gliosis have been greater in BALB/c mice.

In the initial phase of the inflammation (7 and 30 DAI), BALB/c mice presented lesions such as gliosis in the parenchyma adjacent to the lateral ventricles. This may have occurred due to the triggered type of immune response (cellular or humoral), the control of the parasitary growth, the ventricle expansion or the presence of circulating antibodies. Reports from the literature describe that it is not necessary the presence of intraventricular cysts for the development of lesions, such as ventriculomegaly, and even the obstruction of the cerebrospinal fluid flow as the presence of circulating antigens from the parasite are enough to cause those reactions, even without the inflammatory reaction in the cavity\textsuperscript{20-21}. The activation and proliferation of the microglia, which is named microgliosis, were discretely higher in C57BL/6 mice throughout the experiment. These data are in accordance to the cellular profile of response, type 1, observed in these animals\textsuperscript{21}. These cells are the main guard ones of the CNS. Once there is a lesion in the encephalic tissue, the microglia goes through an activation process in which there is the modification of its morphology, surface phenotype characteristics, hypertrophy, increase in the expression of complement receptors such as CR3, increase in the expression of molecules from the main histocompatibility complex (MHC), transforming the cell into one more capable of defending and stimulating the regeneration of the destroyed nervous tissue\textsuperscript{24-25}.

In the late phase of the inflammation (60 and 90 DAI), in spite of BALB/c mice presented a greater intensity in the inflammatory infiltration than C57BL/6 ones, the latter were more capable of containing the growth of the cysticerci and of inducing their death. The inflammatory infiltration with the predominance of polymorphonuclear cells was a remarkable characteristic of the infected BALB/c mice, while in C57BL/6 mice the predominance was of mononuclear cells. Fragoso et al.\textsuperscript{21}, when evaluating the susceptibility and resistance of those both lineages against \textit{T. crassiceps} cysticerci intraperitoneal infections, reported that C57BL/6 mice presented lower parasitary growth and development of lesions and a predominance of type 1 immune response. On the other hand, BALB/c mice presented a predominance of type 2 immune response which is humoral\textsuperscript{26-27}. All infected animals from both lineages from this study presented discrete areas of fibrosis, which demonstrates tissue destruction due to the action of the parasite and the host inflammatory response aiming at eliminating the pathogen and the finalization of the inflammatory process. In the animals from the Control Group, we also observed discrete edema and hyperemia in the initial days of the experiment. These reactions were
not observed in the subsequent days of the experiment and we believe that they occurred due to the sterile saline inoculation procedure.

The dystrophic calcification areas observed in infected C57BL/6 mice at 90 DAI are in accordance to the findings from other authors²²,⁶,²⁸. According to the human NCC classification proposed by Cuéter et al.⁴,⁵, in the intraventricular active form may be an obstruction of the cerebrospinal fluid flow and hypodensic areas in the nuclear magnetic resonance examination, while in the inactive form there is the late hydrocephaly without adjacent calcification areas. Probably the strong type 1 immune response, which is characteristic of the C57BL/6 mice⁶,⁹, may be responsible for the injuries to the adjacent tissues right at the initial phase of the infection, resulting in the contention of the parasitary growth, its death and calcification.

The experimental model presented in this study, which used T. crassiceps cysticerci, may become a reproducible method for human NCC studies because the parasite caused a dynamic inflammatory response from the host, which evolved throughout the experimental days and also because the Control Group presented minor lesions, such as discrete hyperemia and edema, due to the inoculation procedures only in the initial days of the experiment. T. crassiceps cysticerci present other advantages such as its rapid development cycle, facilities in maintenance, and antigenic similarities to T. solium cysticerci²⁵,²⁸.

The previously described NCC model in the literature uses Mesocestoides corti²⁸,¹¹, which are not even from the Taenia genus and do not present the cysticercus evolutive form, they only present the cysticercoid and tetrathyridium forms. Also, the M. corti experimental model presents another drawback as this parasite can proliferate and invade brain tissue¹¹,¹², which was not observed in the T. crassiceps model. The main studies related to the immune and inflammatory responses in experimental cysticercosis caused by T. crassiceps report their findings through the intraperitoneal model⁹. The model proposed in this study may reproduce NCC and the inoculation of the parasite could occur in several areas of the encephalon such as the hippocampus, which may be correlated to seizures that represent the most common human clinic form of NCC.

Therefore, it was possible to induce NCC in both mice lineages proving it to be a good experimental model. The lesions and alterations observed were ventriculomegaly, perivasculitis, meningitis, microgliosis, and inflammation. The observation of the pathological processes in the encephala removed from the infected animals it is possible to conclude that BALB/c mice are less efficient in inducing precocious necrosis of the parasite and they present an acute inflammatory profile. While the C57BL/6 mice are more capable of provoking the parasite’s death, they have a chronic inflammatory profile, less intensity of the alterations and lesions and therefore are considered more resistant to T. crassiceps cysticerci infection.

References

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