NK cells infiltrating in the central nervous system aggravate brain injury of mice caused by Angiostrongylus cantonensis infection

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Research

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Abstract

Background Angiostrongylus cantonensis (A. cantonensis), is a food-borne zoonotic parasite that can cause central nervous system (CNS) injury characterized by eosinophilic meningitis. However, the pathogenesis of the neurological impairments caused by A. cantonensis infection has not been well elucidated. Natural killer cells (NK cells) are unique innate lymphocytes important in early defense against pathogens. It was reported that NK cells could migrate to the CNS after brain injury. The aim of the present study was to investigate the role of NK cells in brain injury caused by A. cantonensis infection.

Methods Mouse model of A. cantonensis infection was established by intragastric administration of third-stage larvae. Neurological impairments were evaluated by Longa's score, Clark's general score and Clark's focal score. Histopathological changes were observed by hematoxylin and eosin staining. The expression of cytokines at gene and protein levels was analyzed by PCR and ELISA, respectively. Infiltration of NK cells in the CNS was detected by immunohistochemistry and flow cytometry. Depletion of NK cells in infected mice was caused by tail vein injection of anti-asialo GM1 rabbit serum, and adoptive transfer of NK cells was performed by tail vein injection of purified splenic NK cells. NK cell-mediated cytotoxicity against YAC-1 cells was detected by LDH release assay. The cytokine production ability was determined by intracellular flow cytometry and ELISA.

Results Mice developed brain inflammation and neurological impairment after A. cantonensis infection. The infiltration of NK cells in the CNS of A. cantonensis-infected mice was observed on 14 dpi and reached the peak on 22 dpi. Compared with the normal splenic NK cells, the CNS-infiltrated NK cells of infected mice expressed lower levels of CD69, NKp46 and NKG2D, but higher levels of NKG2A, and showed enhanced cytotoxicity and increased IFN-γ and TNF-α production ability. Depletion of NK cells alleviated brain injury, whereas adoptive transfer of NK cells exacerbated brain damage in A. cantonensis-infected mice.

Conclusions Our results demonstrate that NK cells infiltrate into the CNS and aggravate the brain damage after A. cantonensis infection. The findings improve the understanding the pathogenesis of angiostrongyliasis and expand the therapeutic intervention in CNS disease.

Background

Angiostrongylus cantonensis (A. cantonensis), is a food-borne zoonotic parasite that can cause damage to the central nervous system (CNS) [1]. It was first discovered in the pulmonary arteries and hearts of domestic rats in Guangzhou, China, by Chen in 1935 [2]. So far, nearly 3000 cases of human angiostrongyliasis have been documented worldwide [3]. With the development of global logistics transportation and climate warming, A. cantonensis has spread from its traditional endemic regions of Southeast Asia and the Pacific islands to the American continent, Europe, Africa and Australia [4–9]. Therefore, angiostrongyliasis has become a potentially fatal globally emerging infectious disease.
Humans and mouse are both non-permissive hosts of *A. cantonensis* and become infected via ingestion of raw or undercooked intermediate hosts including snails or slugs, or vegetables or water contaminated by the infective third-stage larvae [3]. After penetrating the intestinal wall, the larvae migrate in the body with the flow of blood, and finally settle in the CNS. These larvae in non-permissive hosts cannot develop into adults, but can survive in the form of larvae for a long time [10]. Most patients when infected with *A. cantonensis* develop eosinophilic meningitis and common clinical symptoms include headache, fever, neck stiffness, paresthesia and vomiting [11]. However, the pathogenesis of angiostrongyliasis is not fully understood. Mechanical damage to the CNS caused by the larvae’ movements, inflammation and immune response caused by the larval secretion and excreta, may be involved [11–13].

Natural killer cells (NK cells) are a type of cytotoxic lymphocytes critical to the innate immune system [14]. NK cells lack T and B cell receptors and undergo activation without antigen presentation. They are able to rapidly kill target cells by cytotoxicity and produce an abundance of cytokines [15]. NK cells are dispersed throughout lymphoid and non-lymphoid tissues. It was reported that NK cells could migrate to the CNS under several pathological conditions, such as brain ischemia, traumatic injury, or infections [16]. However, very little is known about whether NK cells are involved in the CNS damage caused by *A. cantonensis* infection.

The objective of this study was to investigate the role of NK cells in *A. cantonensis* infection. Our findings will be helpful to further elucidate the pathogenesis of angiostrongyliasis and expand therapeutic intervention in CNS disease.

**Methods**

**Animal experiments**

A total of 300 female BALB/c mice (6–8 weeks old and weighing 18–20 g) were purchased from the Animal Core Facility of Nanjing Medical University (China), maintained in a specific pathogen-free environment, and provided unlimited access to food and water. All experiments were performed in strict compliance with the institutional guidelines and were approved by the Institutional Animal Care and Use Committee of Nanjing Medical University (Approval No. IACUC-1812040). Each mouse was infected with 20 *A. cantonensis* third-stage larvae (L3) by intragastric administration. L3 were isolated from *A. cantonensis*-infected *Biomphalaria glabrata* using the method described previously [13]. The mice were euthanized on 10, 14, 18, 22, and 26 days post-infection (dpi).

**Experimental Grouping**

The experiment was grouped according to different objectives.

To detect the effect of *A. cantonensis* infection on mice and the percentage and number of NK cells, the mice were divided into five groups according to the time of infection with three to twelve mice per group: 0, 10, 14, 18 and 22 dpi.
When detecting the phenotypic and functional changes of NK cells, cells were isolated from mice on 18 dpi and divided into three groups with three to four mice per group: splenic NK cells of uninfected mice (uninfected sNK), splenic NK cells of infected mice (infected sNK) and brain NK cells of infected mice (infected bNK).

In the experiment of NK cell depletion or adoptive transfer, the mice were euthanatized on 18 dpi and divided into four groups with three to twelve mice per group: uninfected group, infected group, NK depleted/transferred group and depleted/transferred control group.

**Neurological Impairment Evaluation**

Longa's score, Clark's general score and Clark's focal score were used to evaluate the neurological impairment of mice infected with *A. cantonensis*. Longa's score is based on a five-point scale, where 0 point indicates no neurologic deficit, 1 point (failure to extend forepaw fully) a mild focal neurologic deficit, 2 point (circling to one side) a moderate focal neurologic deficit, 3 point (falling to one side) a severe focal deficit, and animals with 4 point cannot walk spontaneously or lose consciousness [17]. Clark's general score includes hair, ears, eyes, posture, autonomous movement, and epileptic seizures. Clark's focal score comprises of body symmetry, gait, climbing, rotation test, forelimb symmetry and beard reactivity [18]. Clark's general score and Clark's focal score are between 0 and 28. The higher the score, the more serious the neurological impairment is. The mice were scored by two technicians using a blind method.

**Histopathological Examination**

Mice were perfused transcardially with 0.9% sodium chloride followed by 4% paraformaldehyde after anesthetized with 2% pentobarbital sodium (Sigma-Aldrich, USA). Brain samples were collected, fixed in 10% neutral formalin, embedded in paraffin, and cut into 3 µm-thick sections. Brain sections were then deparaffinized in xylene, rehydrated via graded alcohols and stained with hematoxylin and eosin (H&E) (Biosharp, Wuhan, China). The sections were observed and photographed under a light microscope (Leica, Heidelberg, Germany).

For immunohistochemistry (IHC) analysis, brain sections were subjected to antigen retrieval by boiling the slices in citrate buffer (pH 6.0) with high heat for 15 min. Then sections were treated with 3% H₂O₂ for 10 min to remove endogenous peroxidase, blocked with 5% rabbit serum at room temperature for 20 min, and incubated with rabbit anti-mouse CD49b monoclonal antibody (mAb) (Abcam, Cambridge, UK) at 4 °C overnight. After being washed in PBS, the sections were incubated with an HRP-conjugated secondary antibody (DAKO, Glostrup, Denmark) at room temperature for 15 min and then stained with 3, 3’-diaminobenzidine (DAB) for 10 min. Haematoxylin was used for cell nuclei detection. The sections were visualized and digitally scanned with a light microscope.

**Quantitative Reverse Transcription PCR (QRT-PCR)**
Total RNA was extracted from mouse brains using TRIzol Reagent (Thermo Fisher Scientific, USA) and reverse-transcribed to cDNA using a PimerScript™ RT Master Mix (TaKaRa, Kusatsu, Japan). qRT-PCR was performed on the LightCycler480® Real-Time PCR System (Roche, Reinach, Switzerland) with the RealUniversal Color PreMix (SYBR Green) (Tiangen, Beijing, China), in accordance with the manufacturer's instructions. The primer sequences were shown in Table 1. The mRNA levels of these genes were measured by the Ct value (threshold cycle), and the relative expression levels were calculated with the $2^{-\Delta\Delta Ct}$ method.

### Table 1

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer sequence (5’→3’)</th>
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</tr>
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<td>IL-1β</td>
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<tr>
<td>IL-6</td>
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<td>TNF-α</td>
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<tr>
<td>GAPDH</td>
<td>AGGTCGCTGTAACGGATTTG</td>
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</tr>
</tbody>
</table>

**Detection Of Cytokine Expression In Brain Tissue**

Each brain tissue was added into 4 mL tissue lysate (RayBiotech, USA) and 20 µL protease inhibitor (Merck, Germany) and homogenized in a gentle MACS separator (MiltenyiBiotec, Bergisch Gladbach, Germany). The total protein concentration in each sample was detected by BCA Protein Assay Kit (Tiangen, Beijing, China) according to the manufacturer's instructions. And then, the levels of cytokines (IL-1β, IL-6 and TNF-α) in brain tissue homogenate samples were determined with commercial ELISA Kits (MultiSciences, Hangzhou, China) according to the manufacturer's manuals.

**Cell Isolation**

Blood was collected by eyeball bleeding and gathered in tubes with 1% heparin sodium (Sigma-Aldrich, USA). The mice were anesthetized and perfused as described previously. And then, the brain, spleen, tibia, and femur were collected from the mice respectively.

Brain tissues were homogenized in grinders and filtered through a 70-µm cell strainer. And then cell pellets were resuspended in 30% Percoll (GE Healthcare, Pittsburgh, USA) and centrifuged against 70% Percoll. The cells between the 30–70% Percoll interfaces were collected as the brain mononuclear cells. Spleen was grinded with a syringe core and filtered through a nylon membrane. Erythrocytes were lysed and removed using Red Cell Lysis Buffer (Beyotime, Shanghai, China). Blood was double diluted and layered on the Ficoll-Paque (GE Healthcare, Pittsburgh, USA). After centrifugation, peripheral blood mononuclear cells (PBMCs) were distributed between plasma and Ficoll-Paque. PBMCs were transferred and resuspended in PBS. The tibia and femur bones were used to prepare bone marrow cells. The
The medullary cavity was washed repeatedly with RPMI-1640 (Gibco B, Gaithersburg, MD, USA) by syringe and bone marrow cells were collected and separated from erythrocytes.

NK cells were purified from brain mononuclear cells and splenic lymphocytes using a magnetic cell sorting system (MACS) incorporating anti-mice CD49 MicroBeads (MiltenyiBiotec, Bergisch Gladbach, Germany), following the manufacturer’s instructions. The purity of CD3<sup>-</sup>CD49<sup>+</sup> NK cells after sorting was over 90% detected by FCM (Additional file 1: Figure S1).

**Flow Cytometry (FCM)**

Cells isolated from the brain, spleen, peripheral blood, and bone marrow were prepared to single cell suspension and resuspended in FCM buffer (0.5% BSA in PBS). For Cell surface marker detection, cells were incubated with TruStain FcX™ anti-CD16/32 (Biolegend, San Diego, USA) to block Fc-receptor for 5 min at 4 °C and then stained with the following specific antibodies: anti-CD45-percp-cy5.5, anti-CD3-FITC, anti-CD49b-APC, anti-CD122-PE, anti-CD69-PE, anti-NKp46-PE, anti-NKG2D-PE, anti-NKG2A-PE, anti-CD107a-PE mAbs or isotype controls (Biolegend, San Diego, USA) for 30 min at 4 °C. Cells were then detected on a Verse flow cytometer (BD Biosciences, San Jose, CA, USA). Data analysis was performed using FlowJo software (TreeStar, Ashland, USA).

For intracellular cytokine analysis, cells were cultured at a density of 2 x 10<sup>6</sup>/ml densities in 12-well plates and stimulated with 2 µL/mL Leukocyte Activation Cocktail plus GolgiPlug (BD Biosciences, San Jose, CA, USA) for 5 h. Cells were collected and stained with anti-CD45-percp-cy5.5, anti-CD3-FITC, anti-CD49b-APC mAbs for 30 min at 4 °C. After washing, cells were fixed and permeabilized using Cytofix/Cytoperm™ Fixation/Permeabilization Kit (BD Biosciences, San Jose, CA, USA) according to the manufacturer’s instructions. And then cells were incubated with anti-TNF-α-PE, anti-IFN-γ-BV421 mAbs or isotype controls (Biolegend, San Diego, USA) for 30 min at 4 °C. Cells were detected and data were analyzed as described previously.

**NK Cell Cytotoxicity Assays**

NK cell-mediated cytotoxicity was determined using the Cytotoxicity Detection Kit<sup>PLUS</sup> (Roche, Reinach, Switzerland) based on the measurement of LDH released from damaged cells according to the manufacturer’s manual. Purified NK cells (as effector cells) were incubated with YAC-1 cells (as target cells) at various effector cell/target cell ratios (1:1, 5:1, 10:1, 20:1) in 96-well plates for 3.5 h. All test samples were prepared in triplicate. Reaction mixture and stop solution were added into each well in turn. The absorbance of the samples was measured at 490 nm by an ELISA reader. The percentage of NK cell-mediated cytotoxicity was calculated by ODs using the following equation: Cytotoxicity (%) = (effector/target cell mix - effector cell control - low control) / (high control - low control) x 100.

**Detection Of NK Cells Cytokine Secretion**
Freshly purified NK cells were cultured in 24-well plates at a density of $1 \times 10^6$ /mL in the presence of 1 ng/mL IL-12 (PeproTech, Rocky Hill, NJ, USA). After 24 h of incubation, the culture supernatants were harvested. The concentrations of TNF-α and IFN-γ in supernatants were detected using Mouse TNF-α or IFN-γ High Sensitivity ELISA Kits (MultiSciences, Hangzhou, China) according to the instructions of the manufacturer.

**Depletion Of NK Cells**

5–7 Ganglio-N-tetraosylceramide (asialo GM1) is a glycolipid expressed on NK cells in mice, rats, and humans. Depletion of NK cells in vivo in infected mice was induced by tail vein injection of anti-asialo GM1 rabbit serum (Wako Pure Chemical Industries, Japan) according to the instructions from manufacturer and other researchers [19, 20]. Injection time and dose were shown in Table 2. Normal rabbit serum (Abbkine, California, USA) were given to the infected mice at the same time as control. To evaluate the effect of NK cell depletion, the mice were euthanatized at 18 dpi and the percentage and number of NK cells in brain and spleen were analyzed by FCM.

<table>
<thead>
<tr>
<th>Injection</th>
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<th>3rd</th>
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<tr>
<td>Days post-infection</td>
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<td>5</td>
<td>10</td>
<td>15</td>
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<tr>
<td>Dose (µL per mouse)</td>
<td>20</td>
<td>20</td>
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**Adoptive Transfer Of NK Cells**

The splenic NK cells from normal mice were purified using MACS and were suspended in PBS. The NK cells ($1 \times 10^6$ /mouse in 200 µL PBS) were transferred to the infected mice by tail vein injection on 12 dpi according to the protocols published by other investigators [21]. The respective diluents were injected to the control mice simultaneously. To evaluate the effect of adoptive transfer, the mice were euthanatized at 18 dpi and the percentage and number of NK cells in brain and spleen were analyzed by FCM.

**Statistical analysis**

Statistical analyses were performed using GraphPad Prism 5.0 (GraphPad Software, San Diego, USA). Survival curve comparison was determined using Log-rank Test. Comparison of the neurological impairment scores was performed using non-parametric test. The difference between two groups was compared using independent-samples T test. Multiple comparison procedures were carried out with one-way analysis of variance (ANOVA). The correlation between the percentage and number of NK cells in different tissues was analyzed by linear correlation. The data are presented as the mean ± standard deviation (SD). A $P$ value $< 0.05$ was considered statistically significant.
**Results**

*A. cantonensis* infection causes neurological impairment and brain inflammation in mice

We constructed a mouse model of *A. cantonensis* infection by intragastric administration of third-stage larvae. To evaluate the pathological damage caused by *A. cantonensis* infection, the survival rate, body weight, neurological function, histopathology, and expression of cytokines in brain tissue were detected on 0, 10, 14, 18 and 22 dpi.

The survival rate of mice decreased gradually with the extension of infection time (Fig. 1a). Only 50% of the mice survived on 22 dpi and the survival rate of mice was merely 8.33% on 30 dpi. The weight of infected mice decreased significantly compared with that of uninfected mice on 18 dpi and 22 dpi (Fig. 1b).

After infection, various symptoms of neurologic deficit appeared in mice including erect hair, arched back, falling to one side when walking, and circling to one side continuously when their tails were lifted. Some mice even could not walk spontaneously, became blind or lost consciousness (Fig. 1c). The Longa's score, Clark's general score and Clark's focal score used to evaluate neurological impairment elevated significantly on 18 and 22 dpi compared with that of 0 dpi (Fig. 1d-f).

Then we observed the histopathological changes of brain tissue in mice. On 10 dpi, there were a few red bleeding spots in the brain tissue, and 1–2 brown, slim, and curving fourth-stage larvae (L4) appeared on the surface. The increased hemorrhage and more invasive larvae were found in the brain tissue from 14 to 22 dpi (Fig. 1g). And then, we prepared the brain tissue sections with H&E staining. On 14 dpi, the meninges became thicker and a few inflammatory cells infiltrated under the meninges. On 18 dpi, the number of infiltrating inflammatory cells increased. On 22 dpi, the meninges were damaged, a large number of inflammatory cells infiltrated under the meninges, and multiple areas of red bleeding area were found (Fig. 1h).

We further examined the gene and protein levels of various inflammatory cytokines in brain tissue. The mRNA levels of IL-1β, IL-6, TNF-α and the protein levels of IL-6 increased significantly on 14, 18 and 22 dpi, and the protein levels of IL-1β and TNF-α elevated significantly on 14 and 18 dpi, compared with that on 0 dpi (Fig. 1i-n).

**NK cells infiltrate into the CNS of mice infected with *A. cantonensis***

H&E staining showed that many inflammatory cells infiltrated into the CNS after *A. cantonensis* infection. In order to detect the presence of NK cells in these inflammatory cells, we performed IHC staining on brain tissue sections using anti-CD49b mAb as NK cell marker. Figure 2a, b showed that no NK cell was observed in the CNS on 0 dpi and 10 dpi. On 14 dpi, a small number of NK cells stained brown appeared under the meninges indicating that NK cells began to infiltrate into the CNS. On 18 dpi, more NK cells appeared in the CNS and the number of CNS-infiltrated NK cells peaked on 22 dpi.
To monitor NK cells quantitatively, brain mononuclear cells were isolated and analyzed by FCM at various time points. As shown in Fig. 2c-e, almost no NK cells were detected in the brain tissue at 0 dpi and 10 dpi. NK cells appeared in the CNS from 14 dpi and the percentage and number of CNS-infiltrated NK cells increased gradually with the extension of infection time. The highest percentage of CNS-infiltrated NK cells appeared on 22 dpi (22 dpi vs 0 dpi: 17.47 ± 6.11% vs 0.45 ± 0.12%, P<0.001) and the maximum number of CNS-infiltrated NK cells were found on 18 dpi (18 dpi vs 0 dpi: 1.92 ± 0.43 × 10^5 vs 696.9 ± 617.2, P<0.001).

To elucidate the origin of NK cells in the CNS, we further analyzed the percentage and number of NK cells in the splenic lymphocytes, peripheral blood mononuclear cells and bone marrow cells by FCM. The results showed that the percentage and number of NK cells in splenic lymphocytes decreased on 14, 18 and 22 dpi. (Fig. 3a, d, e). So did the number of NK cells in PBMCs from 10 dpi to 22 dpi (Fig. 3b, e, f). Furthermore, the percentage and number of sNK (NK cells in spleen) were negatively correlated with those of bNK (NK cells in brain) (r =0.79, P<0.01; r =0.85,P<0.01) (Fig. 3g, h). The percentage and number of pbNK (NK cells in PBMCs) and bNK were also negatively correlated, but there was no statistical significance (Fig. 3i, j).

However, the percentage of total NK cells (bmNK, CD122^+), NK precursor cells (NKP, CD49b^-CD122^+) and mature NK cells (mature NK, CD49b^+CD122^+) in bone marrow all significantly increased on 18 and 22 dpi. (Fig. 4a-d). And the ratio of NKP to mature NK in bone marrow up-regulated (Fig. 4e). In addition, the percentage of total NK cells in bone marrow was positively correlated with that of bNK (r = 0.87, P<0.001) (Fig. 4f).

CNS-infiltrated NK cells of A. cantonensis-infected mice have elevated cytotoxicity and secretory ability

We further detected the phenotypic and functional changes of NK cells after A. cantonensis infection. We measured the expression of activation marker CD69, activated receptor NKp46 and NKG2D, and inhibitory receptor NKG2A on NK cells by FCM. Figure 5a-g revealed that infected bNK expressed lower levels of CD69, NKp46 and NKG2D, but higher levels of NKG2A, and infected sNK expressed lower levels of CD69, NKp46, but higher levels of NKG2A, compared with uninfected sNK. Although the phenotypic changes of infected bNK were greater than those of infected sNK, there was no statistical difference between them.

Then we detected NK cell-mediated cytotoxicity against YAC-1 cells by LDH release assay. As shown in Fig. 6a, infected bNK and infected sNK had enhanced cytotoxicity, compared with uninfected sNK (Effect cells: Target cells = 20:1, 19.37% vs 10.59% vs 8.35%). Furthermore, the expression of CD107a, a surface marker of NK cell degranulation, on infected bNK and infected sNK was up-regulated compared with that of uninfected sNK (Fig. 6b, c).

The ability of NK cells to secrete cytokines was determined by two methods: ELISA (secretory levels) and FCM (intracellular levels). Firstly, we detected the concentrations of TNF-α and IFN-γ in the culture supernatant of NK cells after IL-12 stimulation using ELISA. The levels of TNF-α and IFN-γ secreted by infected bNK and infected sNK were significantly higher than those of uninfected sNK (Fig. 6d).
Subsequently, purified NK cells were stimulated and measured by FCM. The percentage of IFN-γ+ cells in infected bNK was higher than that of uninfected sNK, while the percentage of TNF-α+ or IFN-γ+ cells in infected sNK did not change significantly (Fig. 6e, f).

**NK cells aggravate brain injury of mice caused by A. cantonensis infection**

We designed NK cell depletion and adoptive transfer experiments to elucidate the role of NK cells in brain injury caused by *A. cantonensis* infection.

Firstly, NK cells were depleted by tail vein injection of anti-asialo GM1 serum to infected mice. The percentage and number of NK cells in brain and spleen of infected mice significantly decreased after NK cell depletion (Fig. 7a-d). The survival rate of NK-depleted mice increased (*P* < 0.01) (Fig. 7e), body weight elevated (17.35 ± 1.38 g vs 15.61 ± 1.1 g, *P* < 0.05) (Fig. 7f), neurological impairment score decreased slightly (Fig. 7g-i), brain inflammation alleviated (Fig. 7j), and the levels of inflammatory cytokines in brain tissue reduced (IL-1β 25.88 ± 2.38 pg/mg vs 30.75 ± 2.83 pg/mg, IL-6 12.88 ± 1.23 pg/mg vs 15.08 ± 1.13 pg/mg, TNF-α 57.17 ± 4.20 pg/mg vs 64.68 ± 4.34 pg/mg, *P* < 0.05) (Fig. 7k-p), compared with the infected mice on 18dpi. These results showed that depletion of NK cells alleviated brain injury in *A. cantonensis*-infected mice.

And then purified splenic NK cells were transferred to infected mice by tail vein injection. The percentage of NK cells in brain and spleen of infected mice increased significantly after adoptive transferring NK cells (Fig. 8a-d). Compared with the infected mice, the survival rate, body weight and neurological impairment score of NK-transferred mice did not change significantly (Fig. 8e-i), but brain inflammation aggravated (Fig. 8j), and the concentrations of inflammatory cytokines in brain tissue elevated (IL-1β 36.36 ± 1.89 pg/mg vs 30.75 ± 2.83 pg/mg, *P* < 0.05; IL-6 18.08 ± 0.55 pg/mg vs 15.08 ± 1.13 pg/mg, *P* < 0.01; TNF-α 76.84 ± 1.21 pg/mg vs 64.68 ± 4.34 pg/mg, *P* < 0.01) (Fig. 8k-p) on 18dpi. It was suggested that adoptive transfer of NK cells exacerbated brain damage in mice caused by *A. cantonensis*.

**Discussion**

Angiostrongylosis, a food-borne parasitic disease, is caused by the larvae of *A. cantonensis* in the host's central nervous system [22]. NK cells are important innate immune effector cells. They can be swiftly mobilized by danger signals and are among the earliest arrivals at target organs against pathogen infection [23]. However, the role of NK cells in the CNS damage caused by *A. cantonensis* infection remains elusive. Our previous work has reported that NK cells in the spleen and peripheral blood showed quantitative reduction and functional changes in an *A. cantonensis*-infected mice model [24]. In the current study, we focused on the NK cells in the CNS. We found a large number of NK cells infiltrated into the CNS of mice after *A. cantonensis* infection and these CNS-infiltrated NK cells had elevated cytotoxicity and secretory ability. Moreover, we demonstrated that NK cells aggravated brain injury of mice caused by *A. cantonensis* infection.
Mice and humans are both non-permissive hosts of *A. cantonensis*, and the pathogenic process is comparatively similar [25]. Consistent with other reports [26–28], we successfully constructed a mouse model infected with *A. cantonensis* and observed serious neurological damage from 18 dpi to 22 dpi. What is noteworthy in this study is that we confirmed that NK cells infiltrated into the CNS after *A. cantonensis* infection. The results of IHC and FCM showed that NK cells began to appear in the brain tissues at 14 dpi. The percentage and number of NK cells increased gradually with the extension of infection time until 22 dpi. The CNS, including the brain and spinal cord, is considered as an immune privileged organ because of the low permeability of the brain-blood barrier (BBB). However, it is conceivable that peripherally activated lymphocytes, including NK cells, might also be able to penetrate the BBB and infiltrate into the CNS under some pathological conditions. In human ischemic brain tissue and a permanent middle cerebral artery occlusion (pMCAO) mouse model, infiltration of NK cells into the ischemic infarct region are observed [29–31]. NK cells are activated in the periphery and then migrated into the CNS of experimental autoimmune encephalomyelitis (EAE) mice [32]. NK cells can be detected in mouse CNS tissues during a variety of infections, including Semliki Forest virus (SFV) [33], murine coronavirus [34] and *L. monocytogenes* [35]. NK cells are also recruited to the CNS in glioma-bearing mice and constitute approximately 50% of all leukocytes in the CNS [36].

To explore the origin of NK cells in the CNS, we examined the distribution of NK cells in *A. cantonensis*-infected mice. The results showed that the percentage and number of NK cells in spleen and in peripheral blood both decreased, consistent with our previous study [24], while the percentage of NK cells in bone marrow increased after *A. cantonensis* infection. In addition, the percentage and number of splenic NK cells were negatively correlated with those of brain NK cells, while the percentage of NK cells in bone marrow was positively correlated with brain NK cells. It was suggested that the CNS-infiltrated NK cells probably were related with the increased hematopoiesis of bone marrow and migration of peripheral NK cells after *A. cantonensis* infection. NK cells originate from bone marrow, are mainly distributed in peripheral blood and spleen, and some lymphatic tissues [15]. However, the distribution of NK cells is not static because these cells can recirculate between organs. NK cells can respond to a large array of chemokines and be recruited to distinct sites in several pathological circumstances [16]. The detailed trafficking patterns of NK cells are not very well characterized. Nevertheless, it appears that chemokines produced by cells that are unique to specific organs may have a role in orchestrating NK cell migration to each organ [16]. It was reported that NK cells might be recruited to the CNS by chemokines such as CX3CL1 produced by neurons [37] and CCL2 and CXCL10 produced by microglia, astrocytes or infiltrating inflammatory cells [29, 38]. The mechanism of recruitment of NK cells to the CNS after *A. cantonensis* infection needs to be further studied.

NK cells are a kind of special lymphocytes, which can lyse the target cells directly and produce various cytokines to regulate immune response [39]. To further identify the characteristics of the CNS-infiltrated NK cells, we detected the phenotype and function of NK cells after *A. cantonensis* infection. The results showed that the expression of activation molecule CD69, activating receptor NKp46 and NKG2D on the CNS-infiltrated NK cells of infected mice was decreased compared with the splenic NK cells of uninfected
mice, while the expression of inhibitory receptor NKG2A increased. Consistent phenotypic changes were observed in the splenic NK cells of infected mice except for NKG2D. CD69 is one of the early activation markers on NK cells and is involved in cell proliferation and signal transduction. Meanwhile, CD69 is a novel regulator of the immune response, which can inhibit the cytotoxicity of NK cells by inducing the production of TGF-β. It was demonstrated that the administration of anti-CD69 mAbs can activate resting NK cells, resulting in a substantial increase in both NK-cell cytolytic activity and IFN-γ production [40]. NK cells express an array of inhibitory and activating receptors recognizing self-ligands or microbial molecules on infected and tumor cells. Coordinated acquisition of the signals originating from inhibitory and activating receptors regulates the effector functions of NK cells. The down-regulation of CD69 and activating receptors and up-regulation of inhibitory receptors on NK cells following *A. cantonensis* infection might imply the changes of their effector functions.

NK cells have a variety of biological functions, with the most important role being cytotoxicity. NK cells can directly lyse malignant cells or pathogen infected cells by several different mechanisms, including perforin/granzyme, Fas/FasL, TNF-α/TNFRI and other pathways [15]. In our study, we found that the CNS-infiltrated NK cells of infected mice had an enhanced cytotoxicity against YAC-1 cells with higher expression of CD107a. CD107a, also known as Lysosome associated membrane protein-1 (LAMP-1), is a marker for degranulation of NK cells and CD8⁺ T cells [41]. It is reported that CD107a expression correlates with NK cell-mediated lysis of target cells and is a sensitive marker of NK cell activity [42]. NK cells can also produce a variety of cytokines in response to activation signaling to mediate immunomodulatory effects. Our results showed that the CNS-infiltrated NK cells of infected mice produced higher levels of TNF-α and IFN-γ than splenic NK cells of uninfected mice. TNF-α is a proinflammatory cytokine that is not only cytotoxic to tumor cells but also involved in mediating anti-pathogen immune responses. IFN-γ is an important immunomodulator with the function of activating macrophages, promoting MHC molecular expression and antigen presentation, and plays an important role in defending viral, bacterial and parasitic infections [43, 44]. Taken together, the CNS-infiltrated NK cells in *A. cantonensis*-infected mice showed stronger activity with enhanced cytotoxicity and elevated production of TNF-α and IFN-γ. There are no NK cells in the steady-state CNS, but NK cells might migrate into the CNS under certain pathological conditions such as *A. cantonensis* infection. After homing to the inflamed CNS, NK cells become receptive to an array of cellular components that they have not encountered in the periphery. These include astrocytes, microglia, neurons, and eosinophils, which release numerous soluble factors with diversified and perhaps coordinated effects on NK cells [16]. The exact cellular and molecular interactions that shape the phenotype and function of NK cells in the CNS still need to be determined.

We then investigated the role of NK cells in the brain damage caused by *A. cantonensis* infection using NK cell depletion and adoptive transfer experiments. The changes in NK cell percentage and number confirmed that NK cell depletion and adoptive transfer were effective. Our results showed that the survival rate and body weight increased, nerve injury and brain inflammation decreased after NK depletion in *A. cantonensis*-infected mice. On the contrary, the inflammation in brain aggravated after adoptive transfer
of NK cells. We speculated that the CNS-infiltrated NK cells might play a role in exacerbating brain injury after *A. cantonensis* infection. Many studies have reported that NK cells can rapidly accumulated into the CNS under pathological conditions. However, the role of NK cells in brain injury diseases remains complex and paradoxical. Zhiguo Li [31] identified NK cells infiltrates into the CNS during early stages of intracerebral hemorrhage (ICH). These infiltrating NK cells exacerbate BBB disruption and brain edema via cytotoxicity toward cerebral endothelial cells and recruitment of neutrophils. Gan Y [30] reported NK cells mediate exacerbation of brain infarction after ischemia via the disruption of NK cell tolerance, augmenting local inflammation and neuronal hyperactivity. Alsharifi M [33] found that NK cells exert both disease-exacerbating and protective effects in Semliki Forest virus infection in the CNS of mice. The cytolytic activity of NK cells is detrimental, while IFN-γ production is beneficial for recovery from SFV infection. However, Hao J [32] demonstrated that the CNS-resident NK cells have a protective role in the brain of EAE mice, as they inhibit the activation of autoimmune T cells through the killing of activated microglia. Jiang W [45] disclosed that acetylcholine-producing NK cells attenuate CNS inflammation of EAE model via modulation of infiltrating monocytes/macrophages. NK cells play different roles in brain injury, which may be related to the initial factors of the primary disease, the time of immune response, and the overall inflammatory process [30]. After brain injury, brain-intrinsic cells, including microglia, astrocytes and neurons, secrete various chemokines which recruiting a series of inflammatory cells. As an important type of immune cell, peripheral resident NK cells are activated and migrate to brain tissue [16]. NK cells infiltrating into the CNS can build a bridge between immune system and nervous system, which not only produce cytotoxicity or cytolysis to pathogens, but also interact with other cells to affect the progress of brain injury. In the process of *A. cantonensis* infection, NK cells infiltrating in brain tissue have stronger killing function and secrete more IFN-γ and TNF-α. On the one hand, these CNS-infiltrated NK cells were beneficial to clearance of pathogens, on the other hand, they were also involved in the death of neural cells and aggravated brain damage.

Currently, the treatment for angiostrongyliasis includes supportive treatment and corticosteroid therapy [46]. The use of anthelmintic drugs, such as albendazole and mebendazole, to kill worms remains controversial. Because the dead worm lysis in the CNS might cause severe inflammatory response and further damage [10]. Meanwhile, Patients under the treatment of high-dose corticosteroids would experience immune suppression [47]. Therefore, it is urgent to develop new therapeutic interventions for angiostrongyliasis. We hypothesized that selective reduction of NK cell infiltration into the CNS or control their activation may help to alleviate the brain injury caused by *A. cantonensis* infection, which need to be further investigated in our future work.

**Conclusion**

In summary, our study demonstrates that NK cells infiltrate into the CNS of *A. cantonensis*-infected mice. These CNS-infiltrated NK cells display enhanced cytotoxicity and secretary ability, which may contribute to aggravate brain damage. Our findings not only enrich the understanding of *A. cantonensis* pathogenesis but also provide a clue to novel potential therapeutic strategies against CNS disease.
Abbreviations

A. cantonensis: Angiostrongylus cantonensis; BBB: Brain-blood Barrier; bNK: NK cells in brain; bmNK: NK cells in bone marrow; CNS: Central nervous system; DAB: diaminobenzidine; EAE: Experimental autoimmune encephalomyelitis; H&E: Hematoxylin and eosin staining; ICH: intracerebral hemorrhage; IHC: Immunohistochemistry; L3: Third-stage larvae; LAMP-1: Lysosome associated membrane protein-1; MACS: Magnetic cell sorting system; NK cells: Natural killer cells; NKP: NK precursor cells; pbNK: NK cells in peripheral blood; PBMCs: Peripheral blood mononuclear cells; pMCAO: permanent middle cerebral artery occlusion; SFV; Semliki Forest virus; sNK; NK cells in spleen.

Declarations

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Authors’ contributions

ALC conceived the project and designed the experiments. YW designed the experiments, supervised the project, and was involved in all aspects of the submission. RZ performed most of the experiments, analyzed data, and wrote the manuscript. TTM was responsible for the infection and feeding of animals, performed cell isolation and detection by FCM. MQ, CSZ performed detection of cytokine by qRT-PCR and ELISA. CCZ and WW participated in neurological impairment evaluation and NK cell cytotoxicity assays. XJL and YC performed histopathological examination and data analysis. All authors read and approved the final manuscript.

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Availability of data and materials
The datasets used and analyzed during the current study are available from the corresponding author on reasonable request.

**Ethics approval and consent to participate**

All experiments were performed in strict compliance with the institutional guidelines and were approved by the Institutional Animal Care and Use Committee of Nanjing Medical University (Approval No. IACUC-1812040).

**Consent for publication**

Not applicable.

**Competing interests**

The authors declare that they have no competing interests.

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**References**


Figures
A. cantonensis infection caused neurological impairment and brain inflammation in mice. Mice were infected with 20 A. cantonensis third-stage larvae by intragastric administration and monitored at 0, 10, 14, 18 and 22 dpi, respectively. a, b Survival rate and body weight of mice at different infection time points. c Representative images of infected mice with neurologic deficit symptoms. Infected mice displayed various neurologic deficit symptoms including erect hair, arched back, falling to one side when
walking, blindness and so on. d-f Neurological impairment scores of infected mice. Longa’s score, Clark’s general score and Clark’s focal score were used to evaluate the neurological impairment. g Representative images of brain tissue in infected mice. The brains tissue of infected mice displayed obvious hemorrhage and fourth-stage larvae (blue arrow) from 14 to 22 dpi. h Representative histopathological sections of brain tissue in infected mice with H&E staining. The meninges of infected mice became thick and inflammatory cells infiltrated under the meninges (yellow arrow) from 14 to 22 dpi. Images are shown at 100 × magnification (Scale bar, 100 µm). i-n Expression of inflammatory cytokines IL-1β, IL-6 and TNF-α in brain at different infection time points. The gene and protein levels of inflammatory cytokines in the brain of infected mice were detected by qRT-PCR and ELISA. Data are expressed as the means ± SD. Data showed represent analysis from two independent experiments with four to twelve mice per group. Survival curve comparison was determined by Log-rank Test. Comparison of body weight was carried by independent-samples T test between infected and uninfected mice. Comparison of the neurological impairment scores was compared by non-parametric test. Multiple comparisons of gene and protein levels of cytokines at different infection time-points were performed using one-way ANOVA. *P < 0.05; **P < 0.01; ***P < 0.001
Figure 2

NK cells infiltrated into the CNS after A. cantonensis infection. a, b Representative histopathological sections of brain tissue in infected mice with IHC staining. Brain sections were incubated with rabbit anti-mouse CD49b monoclonal antibody and stained with DAB. There were no NK cells in the CNS on 0 dpi and 10 dpi. From 14 dpi to 22 dpi, more and more NK cells (stained brown) appeared under the meninges. Images are shown at 200 × and 400× magnification (Scale bar, 100 µm). c-e The percentage and number of NK cells in brain mononuclear cells of infected mice. NK cells in brain mononuclear cells were detected by FCM. The gating strategy for the mouse CD45+CD3-CD49b+ NK cell population in brain was shown in Additional file 2: Figure S2. Data are expressed as the means ± SD. Data showed represent analysis from two independent experiments with three mice per group. Significance was determined by one-way ANOVA. **P < 0.01; ***P < 0.001
Figure 3

The percentage and number of NK cells in spleen and peripheral blood decreased after A. cantonensis infection. a, c, d The percentage and number of NK cells in splenic lymphocytes of infected mice. b, e, f The percentage and number of NK cells peripheral blood mononuclear cells of infected mice. NK cells in the splenic lymphocytes and peripheral blood mononuclear cells were detected by FCM. g, h The correlation between the percentage and number of NK cells in spleen and brain. i, j The correlation between the percentage and number of NK cells in peripheral blood and brain. Data are expressed as the means ± SD. Data showed represent analysis from two independent experiments with three mice per group. Multiple comparisons of the percentage and count of NK cells at different time-points of infection were performed by one-way ANOVA. The correlation between the percentage and number of NK cells in
different tissues was analyzed by linear correlation. *P < 0.05; **P < 0.01; ***P < 0.001. sNK, NK cells in spleen; bNK, NK cells in brain; pbNK: NK cells in peripheral blood.

Figure 4

The percentage of NK cells in bone marrow increased after A. cantonensis infection. a-d The percentage of NK cells in bone marrow lymphocytes of A. cantonensis-infected mice. NK cells in bone marrow (CD122+) were divided into NK precursor cells (CD122+CD49b-) and mature NK cells (CD122+ CD49b+) detected by FCM. e The ratio of NK precursor cells to mature NK cells in bone marrow. f The correlation between the percentage of NK cells in bone marrow and brain. Data are expressed as the means ± SD. Data showed represent analysis from two independent experiments with three mice per group. Multiple comparisons of the percentage of NK cells at different time-points of infection were performed by one-way ANOVA. The ratio of different time points was analyzed by nonparametric test. The correlation between the percentage and number of NK cells in different tissues was analyzed by linear correlation. **P < 0.01; ***P < 0.001. bmNK, NK cells in bone marrow; NKP, NK precursor cells; mature NK.
Figure 5

The phenotypes of NK cells changed after A. cantonensis infection. The expression of activation marker CD69 (a, b), activated receptor NKp46 (c, d) and NKG2D (e, f), and inhibitory receptor NKG2A (g, h) on the surface of sNK cells and bNK cells were detected by FCM. Data are expressed as the means ± SD. Data showed represent analysis from two independent experiments with three mice per group. Multiple
comparisons of phenotypes between uninfected sNK, infected sNK and infected bNK were performed by ANOVA. *P < 0.05; ***P < 0.001. sNK; NK cells in spleen; bNK, NK cells in brain.

**Figure 6**

The cytotoxicity and secretory ability of NK cells elevated after A. cantonensis infection. a NK cell-mediated cytotoxicity against YAC-1 cells. Purified sNK cells and bNK cells (as effector cells) were incubated with YAC-1 cells (as target cells) at various effector cell/target cell ratios (E:T = 1:1, 5:1, 10:1, 20:1). NK cell-mediated cytotoxicity was detected by LDH release assay and calculated by ODs using the following equation: Cytotoxicity (%) = (effector/target cell mix - effector cell control - low control) / (high control - low control) × 100. b, c The expression of CD107a on NK cells. Splenic lymphocytes and brain mononuclear cells were isolated and detected by FCM. CD45+CD3-CD49b+ NK cells were gated and analyzed for the expression of CD107a. d The concentrations of TNF-α and IFN-γ in the culture supernatants of NK cells. Purified sNKs and bNKs were stimulated with IL-12 and the concentrations of TNF-α and IFN-γ in the culture supernatants were detected using ELISA. e, f The expression of intracellular cytokine TNF-α and IFN-γ in NK cells. Splenic lymphocytes and brain mononuclear cells were isolated and stimulated with Leukocyte Activation Cocktail. The percentage of TNF-α+ cells and IFN-γ+ cells in
CD45+CD3-CD49b+ NK cells was detected by FCM. Data are expressed as the means ± SD. Data showed represent analysis from two independent experiments with four mice per group. Multiple comparisons were performed by one-way ANOVA. *P < 0.05; **P < 0.01; ***P < 0.001. sNK; NK cells in spleen; bNK, NK cells in brain.

**Figure 7**

- **A** and **B**: Bar graphs showing the percentage of NK cells in brain before and after infection and treatment.
- **C** and **D**: Graphs showing the count of NK cells in spleen.
- **E**: Graph illustrating the survival proportion over time.
- **G** to **I**: Graphs showing the scores of Longa's, Clark's general, and focal scores.
- **J**: Images showing HE stain at 100X, IHC at 200X and 400X.
- **K** to **P**: Graphs showing the concentration of IL-1β, IL-6, and TNF-α.

*Note: The text above is a natural representation of the document content.*
Depletion of NK cells alleviated brain injury in A. cantonensis-infected mice. Depletion of NK cells in infected mice was induced by tail vein injection of anti-asialo GM1 rabbit serum. The percentage and number of NK cells in brain (a, b) and spleen (c, d) were detected by FCM to evaluate NK cell depleting efficiency. After NK cell depletion, the survival rate of infected mice (e) increased, body weight (f) elevated, while neurological impairment scores (g-i) did not have significant changes. Representative histopathological sections of brain tissue with H&E and IHC staining (j) showed alleviated inflammation and fewer infiltrating NK cells in the brain of NK-depleted mice. Images are shown at 100 ×, 200 × and 400 × magnification (Scale bar, 50-100 µm). The gene (k-m) and protein levels (n-p) of inflammatory cytokines IL-1β, IL-6 and TNF-α in brain tissue reduced after NK cell depletion measured by qRT-PCR and ELISA. Data are expressed as the means ± SD. Data showed represent analysis from two independent experiments with three to twelve mice per group. Survival curve comparison was determined by Log-rank Test. Comparison of the neurological impairment scores was compared by non-parametric test. Multiple comparisons of the percentage and number of NK cells, body weight and expression of cytokines were performed using one-way ANOVA. *P < 0.05; **P < 0.01; ***P < 0.001. Uninfected, normal mice; Infected, mice infected with A. cantonensis; NK Depleted, infected mice depleted NK cells by injection of anti-asialo GM1 rabbit serum; Depleted Control, infected mice given normal rabbit serum as control.
Figure 8

Adoptive transfer of NK cells exacerbated brain damage in A. cantonensis-infected mice. Purified splenic NK cells from normal mice were transferred to infected mice by tail vein injection. The percentage and number of NK cells in brain (a, b) and spleen (c, d) were detected by FCM to evaluate NK cell adoptive transferring efficiency. After adoptive transferring NK cells, the survival rate of infected mice (e), body weight (f) and neurological impairment score (g-i) did not change significantly. Representative
Histopathological sections of brain tissue with H&E and IHC staining (j) showed aggravated inflammation and more infiltrating NK cells in the brain of NK-transferred mice. Images are shown at 100 ×, 200 × and 400 × magnification (Scale bar, 50-100 µm). The gene (k-m) and protein levels (n-p) of inflammatory cytokines IL-1β, IL-6 and TNF-α in brain tissue elevated after NK cell adoptive transferring measured by qRT-PCR and ELISA. Data are expressed as the means ± SD. Data showed represent analysis from two independent experiments with three to twelve mice per group. Survival curve comparison was determined by Log-rank Test. Comparison of the neurological impairment scores was compared by non-parametric test. Multiple comparisons of the percentage and number of NK cells, body weight and expression of cytokines were performed using one-way ANOVA. *P < 0.05; **P < 0.01; ***P < 0.001. Uninfected, normal mice; Infected, mice infected with A. cantonensis; NK Transferred, infected mice transferred NK cells by tail vein injection; Transferred Control, infected mice injected with PBS as control.

**Supplementary Files**

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