Research Article

Substance P Signaling Contributes to Granuloma Formation in Taenia crassiceps Infection, a Murine Model of Cysticercosis

Armandina Garza,1 David J. Tewardy,1 Joel Weinstock,2 Balaji Viswanathan,1 and Prema Robinson1

1 Section of Infectious Disease, Department of Medicine, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
2 Division of Gastroenterology, Tufts New England Medical Center, 750 Washington St., Box 233, Boston, MA 02111, USA

Correspondence should be addressed to Prema Robinson, premar@bcm.tmc.edu

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Cysticercosis is an infection with larval cysts of the cestode Taenia solium. Through pathways that are incompletely understood, dying parasites initiate a granulomatous reaction that, in the brain, causes seizures. Substance P (SP), a neuropeptide involved in pain-transmission, contributes to inflammation and previously was detected in granulomas associated with dead T. crassiceps cysts.

To determine if SP contributes to granuloma formation, we measured granuloma-size and levels of IL-1β, TNF-α, and IL-6 within granulomas in T. crassiceps-infected wild type (WT) mice and mice deficient in SP-precursor (SPP) or the SP-receptor (neurokinin 1, NK1). Granuloma volumes of infected SPP- and NK1-knockout mice were reduced by 31 and 36%, respectively, compared to WT mice (P < .05 for both) and produced up to 5-fold less IL-1β, TNF-α, and IL-6 protein. Thus, SP signaling contributes to granuloma development and proinflammatory cytokine production in T. crassiceps infection and suggests a potential role for this mediator in human cystercercosis.

1. Introduction

Neurocysticercosis is the most common parasitic disease of the central nervous system leading to seizures worldwide [1]. Humans develop cysticercosis when they ingest eggs of the tapeworm Taenia solium usually found in fecal-contaminated water or food [2]. Neurocysticercosis (NCC) is endemic to many parts of the world [3–7] and is becoming an increasingly important cause of seizures in the United States due to immigration from Mexico and Central and South America [8, 9]. Seizures in NCC most commonly arise as a result of the granulomatous responses to dead cysts in the brain. The granulomatous response is associated with production of several cytokines including T helper 1 (Th1) cytokines such as interferon gamma (IFN-γ), interleukin-2 (IL-2), and interleukin-12 (IL-12) [10].

T. crassiceps infection in mice is an experimental model for T. solium cysticercosis in man [11–15]. Intraperitoneal inoculation with 10 cysts of T. crassiceps results in the entire peritoneal cavity of the mouse demonstrating granulomatous inflammation within 3–6 months. Similar to human infection, minimal granulomatous inflammation is found surrounding live parasites; rather, granulomatous inflammation is initiated when the parasite dies. The mediators contributing to development of granulomas around the dead parasite and production of proinflammatory cytokines are not completely understood.

We previously detected substance P (SP) protein within granulomas associated with T. crassiceps infection [16, 17]. We also demonstrated that levels of IL-2, IFN-γ, IL-4, and IL-10 protein were significantly higher in granulomas from infected WT mice than granulomas from infected SPP-knockout or the SP-receptor (neurokinin 1, NK1) NK1-knockout mice [16, 17]. In addition, we detected mRNA for IL-1α, IL-1β, IL-1 receptor antagonist, and TNF-α in all granulomas derived from infected WT mice [18]. However, corresponding proteins levels were not assessed nor was the contribution of SP signaling to their mRNA and protein production and to granuloma formation.
The current studies were aimed at determining if SP and NK1 contributed to granuloma development and/or to production of IL-1β, TNF-α, and IL-6 in cystercerosis. SP stimulates production of proinflammatory cytokines such as of IL-1β, IL-6, and TNF-α by human peripheral mononuclear cells, bronchial cells, and astrocytes [19–29]. SP also contributes to inflammatory processes associated with other infectious diseases. For example, granulomatous inflammation in murine schistosomiasis requires binding of SP to NK1 [30]. SP has been demonstrated to stimulate inflammatory cell infiltration. SP injection induced recruitment of leucocytes into the pleural cavity of mice and into the skin of humans [19–29] and stimulated the migration of human fibroblasts and peripheral blood lymphocytes in studies using modified Boyden chambers or micropore filter analysis, respectively [19–29]. In the current studies, we determined granuloma size and measured levels of IL-1β, TNF-α, and IL-6 protein within granuloma obtained from *T. crassiceps*-infected WT and mice deficient in SPP or NK1. These studies indicate that SP signaling contributes to granuloma development and proinflammatory cytokine production in *T. crassiceps* infection and suggests a potential role for this mediator in human cystercerosis.

### 2. Materials and Methods

#### 2.1. Murine Cystercrosis Model

Female mice were infected by intraperitoneal inoculation with 10 cysts of the ORF strain of *T. crassiceps*, as described in [16, 17]. Three months following infection, the mice were euthanized by cervical dislocation under anesthesia using a combination anesthetic sacrifice rodent cocktail ketamine, 25 mg/kg, acepromazine 0.8 mg/kg, xylazine 5 mg/kg intraperitoneally, at a dose of 0.5–0.7 mL/kg intramuscularly. Granulomas associated with dying cysts were removed from the peritoneal cavity of each of the infected mice that were euthanized. Three groups of mice were included in the experiments: (1) wild type C57BL/6 mice; (2) preprotachykinin or SPP-knockout mice (Jackson Laboratories, Maine, USA, bred >10 generations onto the C57BL/6 background); (3) NK1-knockout mice provided by Dr. Joel Weinstock, Tufts New England Medical Center, Boston, USA, bred >10 generations onto the C57BL/6 background. Three to 8 infected mice from each of the 3 groups were used for this study; 4–15 granulomas per mouse group were used for this study. Granulomas associated with parasites were identified visually, removed from the peritoneal cavity, and either used for quantifying cytokine proteins by ELISA or used for size determinations. This study was approved by the Animal Research Committee at Baylor College of Medicine.

#### 2.2. Granuloma Size Determination

Intact granuloma was obtained from *T. crassiceps*-infected WT mice (2 granulomas), SPP-knockout mice (4 granulomas), and NK1-knockout mice (3 granulomas), fixed with 4% paraformaldehyde, paraffin imbedded and completely sectioned by microtome into 7 micron sections. Each section was stained with giemsa and examined microscopically. The area of granuloma within each section was measured using Image J software (NIH). The volume of granuloma within each section was calculated by multiplying the area times 7 microns and the volume of granuloma within each section totaled to give the total granuloma volume.

### 2.3. Sandwich ELISA for the Detection of Cytokines

Cytokine protein levels were determined in 12–15 granulomas derived from *T. crassiceps* infected WT mice, 4–6 granulomas derived from *T. crassiceps* infected SPP-knockout mice, and 9–10 granulomas derived from *T. crassiceps*-infected NK1-knockout mice. A portion of each granuloma was homogenized in PBS, followed by centrifugation at 16,000 g. Total protein in the supernatant was quantitated using the Bradford method (cat no. 500-0006, Bio-Rad, Hercules, CA). IL-1β, IL-6, and TNF-α protein levels were determined by sandwich ELISA (R&D Systems, San Diego, California) as per manufacturer’s instruction. Results are expressed as pg cytokine/mg total protein.

### 2.4. Statistical Analysis

Differences between groups were compared using Student’s *t*-test. Significance was set at *P* < .05.

### 3. Results

#### 3.1. Granulomas from *T. crassiceps*-Infected SPP-Knockout and NK1-Knockout Mice Are Smaller than Granulomas Derived from Infected WT Mice

To begin to examine the contribution of SP signaling to granuloma formation in NCC, we measured granuloma volume in mice with normal and deficient SP signaling. The volumes of granulomas from *Taenia crassiceps* infected SPP-knockout mice (1.8 ± 0.45 mm³) and NK1-knockout mice (1.68 ± 0.40 mm³) were reduced by 31% and 36%, respectively, compared to granulomas derived from infected WT mice (2.62 ± 0.28 mm³; *P* < .05 for both; Student’s *t*-test; see Figure 1).

#### 3.2. IL-1β Protein Levels Are Reduced within Granulomas from *T. crassiceps* Infected SPP-Knockout and Infected NK1-Knockout Mice Compared to Granulomas from Infected WT Mice

IL-1β is the primary mediator of granuloma formation in the *S. mansoni* pulmonary granuloma model [31]. Also, intratracheal injection of agarose beads coupled to recombinant IL-1β induced pulmonary granulomas in mice [32]. To determine if decreased production of IL-1β in SPP- and NK1-knockout mice contributed to reduced granuloma size in these animals, we measured IL-1β protein levels in the granulomas derived from each group of mice. IL-1β protein levels in granulomas from *T. crassiceps* infected SPP-knockout mice (216 ± 129 ng/mg total protein) and NK1-knockout mice (101 ± 43 ng/mg total protein) were reduced by 48% and 76%, respectively, compared to granulomas from infected WT mice (418 ± 278 ng/mg total protein; *P* < .05 for both; Student’s *t*-test; see Figure 2). Thus, SP signaling contributes to IL-1β production within granulomas formed in response to dying *T. crassiceps* cysts. Furthermore, reduced levels of this cytokine likely contributed to reduced granuloma size in SPP- and NK1-knockout mice.
3.3. **TNF-α Protein Levels Are Decreased within Granulomas from T. crassiceps Infected SPP-Knockout and Infected NK1-Knockout Mice Compared to Granulomas from Infected WT Mice.** TNF-α is responsible for granuloma development in multiple settings. Intratracheal injection of agarose beads coupled to TNF-α induced pulmonary granulomas in mice [32]. TNF-α mediates granuloma growth in the *S. mansoni* pulmonary granuloma model [31] and is required for granuloma formation in a mouse model of tuberculosis [33].

Similar to our findings with IL-1β, TNF-α protein levels in granulomas from *T. crassiceps* infected SPP-knockout mice (96±67 ng/mg total protein) were reduced by 79% compared to levels in granulomas derived from infected WT mice (460 ± 452 ng/mg total protein; P < .05; Student’s t-test; see Figure 3). TNF-α protein levels within granulomas from NK1-knockout mice (345 ± 153 ng/mg total protein) were decreased by 25% compared to levels with granulomas of WT mice; however, this difference did not achieve statistical significance. Thus, SP contributes to TNF-production within granulomas formed in response to dying *T. crassiceps* cysts. Furthermore, reduced levels of this cytokine likely contributed to reduced granuloma size in SPP- and, perhaps, NK1-knockout mice. The failure to detect a significant difference in TNF-α protein levels between NK1-knockout and WT mice suggests the possibility that SP-mediated increases in TNF-α may occur through binding of SP to other members of the NK family, for example, NK2 or NK3.

3.4. **IL-6 Levels Are Decreased within Granulomas from T. crassiceps Infected SPP-Knockout and Infected NK1-Knockout Mice Compared to Granulomas from Infected WT Mice.** IL-6 production in human peripheral blood mononuclear cells, bronchial cells, and astrocytes is increased directly by SP through the action of nuclear factor IL-6 (NF-IL-6) and p38 MAPK, as well as indirectly in response to IL-1β and TNF-α through the activation of NF-κB [26, 27, 34, 35]. IL-6 mediates its acute proinflammatory effects within infected or injured tissues, in part, through upregulation of CXC chemokines, which leads to recruitment of the first wave of inflammatory cells. As we expected from the IL-1β and TNF-α results summarized above, IL-6 protein levels in the granulomas derived from infected SPP-knockout mice (28 ± 30 ng/mg total protein) and from infected NK1-knockout mice (50 ± 16 ng/mg total protein) were reduced by 79 and 62%, respectively, compared to levels in granulomas from infected WT mice (130 ± 153 ng/mg total protein; P < .05 for
both; Student’s t-test; see Figure 4). Thus, SP signaling either directly or indirectly through the actions of IL-1β and TNF-α contributes to IL-6 production within granulomas formed in response to dying T. crassiceps cysts.

4. Discussion

The current studies were performed to determine the contribution of SP and its specific receptor, NK1, to granuloma development and proinflammatory cytokine production within granulomas arising in mice infected with T. crassiceps. We demonstrated that the size of granulomas from the T. crassiceps-infected SPP-knockout mice and infected NK1-knockout mice were significantly smaller than granulomas from the infected WT mice. Furthermore, proteins levels of IL-1β, a key mediator of granuloma formation, were significantly lower within granuloma from SPP- and NK1-knockout mice compared to granuloma from mice. In addition, compared to granulomas from WT mice, protein levels of TNF-α, another key mediator of granuloma formation, were significantly lower in SPP-knockout mice and trended in the same direction in NK1-knockout mice. Thus, SP signaling contributes to granuloma formation, in part, through induction of IL-1β and TNF-α, key mediators of granuloma formation.

Substance P and its high affinity receptor, NK1, are known to play an important role in inflammatory responses. Granuloma formation in response to murine schistosomiasis requires binding of SP to its specific receptor [30]. SP is known to stimulate inflammatory cell infiltration and to induce the production of proinflammatory cytokines such as IL-1β, IL-6, and TNF-α by human peripheral mononuclear cells, bronchial cells, or astrocytoma cells [19–29]. Our findings extend these observations and indicate that SP signaling contributes to granuloma formation and production of IL-1β, TNF-α, and IL-6 protein within granulomas formed in response to T. crassiceps infection. The mechanism by which SP stimulates the production of these cytokines may be by mediating inflammatory cell influx [19–24]. Nerves, endothelial cells, and cells of the immune system produce SP [36–38]. All of these cells have receptors for SP and are known to respond to SP [38–40]. SP is known to stimulate influx of lymphocytes, monocytes, macrophages, and other immune cells that produce proinflammatory cytokines such as IL-1β, IL-6, and TNF-α [19–29]. Although there are various studies on the molecular mechanisms by which SP stimulates the production of IL-6 and TNF-α, there is limited information on the molecular mechanisms by which SP stimulates the production of IL-1β. Therefore, our results in SPP-knockout mice can be attributed to a deficiency in either SP or neurokinin A or both. However, since NK1 binds only SP and not neurokinin A and the results in NK1-knockout mice mirror the findings in SP-knockout mice, we are confident in attributing reduced granuloma formation and proinflammatory cytokine production to the absence of SP signaling and not to reduced or absent neurokinin A signaling.

In the current studies, we demonstrated that the granuloma size and the levels of the proinflammatory cytokine, IL-1β, are lower in the infected NK1-knockout mice compared to those of the infected SPP-knockout mice. Besides SP, peptide hormones such as hemokinin can also bind and activate NK1 at sites of chronic inflammation [44]. Therefore, although the current studies suggest that SP may be an important mediator associated with cytokine production, there may be other peptide hormones like hemokinin that also bind and activate the NK1 receptor that may be associated with granuloma and cytokine production. Therefore, it may be possible that in the NK1-knockout mice, the synergistic lack of activity of both SP and hemokinin may have resulted in lower IL-1β levels and granuloma size as compared to SPP-knockout mice.

Granuloma formation by the host in response to agents causing chronic infections is thought to be essential for limiting and eventually clearing infection. However, recent work in zebra fish infected with Mycobacterium marinum suggests that granuloma formation contributes to early bacterial growth [45]. Intraparenchymal cysts of NCC are thought to die spontaneously and to elicit a granulomatous response that does not in itself contribute to the demise of...
the cyst. Rather, we have previously demonstrated that early granuloma formed in response to dying cysts contributes to NCC disease manifestations by producing mediators that induce seizures [46]. Other groups have demonstrated that SP is epileptogenic [47]. The current studies demonstrating that SP signaling contributes to granuloma formation in *Taenia crassiceps* infection, together with other published observations, suggest the possibility that diminishing granuloma formation in NCC by blocking SP, which contributes to granuloma formation and epileptogenic responses, may be beneficial in the treatment of this disease.

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**References**


