Expression of TNF-α and IL-1β in Splenic Dendritic Cells and Their Serum Levels in Mouse Sparganosis

Hyun-Jong Yang*
Department of Parasitology and Ewha Global Challenge, School of Medicine, Ewha Womans University, Seoul 158-710, Korea

Abstract: Sparganosis is a tissue invading helminthiasis infecting intermediate hosts, including humans. Strong immune responses are expected to occur in early phases of infection. Thus, we investigated cytokine expressions in splenic dendritic cells and in sera after experimental infection of mice. In splenic dendritic cells, TNF-α and IL-1β expression peaked at week 1 and week 3 post-infection (PI), respectively, and also early phase (week 2 PI) depressed cytokine expression was noticed. Serum IL-1β concentration increased significantly at week 2 PI and peaked at week 6 PI, and that of TNF-α peaked at week 6 PI. These results showed that pro-inflammatory cytokines, TNF-α and IL-1β, are chronologically regulated in mouse sparganosis.

Key words: Spirometra mansoni, sparganum, dendritic cell, TNF-α, IL-1β

After oral infection, the sparganum (the plerocercoid of *Spirometra mansoni*) penetrates the intestinal wall and migrates to the peritoneal cavity within a few hours. Finally, worms move to subcutaneous tissues and form a subcutaneous mass in the skin [1].

Dendritic cells (DC) are known as the most potent antigen-presenting cells due to their ability of inducing primary immune responses and activating immune memory cells [2]. Immature DC in the peripheral tissues capture antigens and have the unique migrating capacity to the T cell areas of secondary lymphoid organs. As the DC migrate, DC alter their expression profiles of cell surface molecules, including chemokine receptors and finally activate the resting T cells in the lymph nodes [3,4].

Pro-inflammatory cytokines, such as TNF-α and IL-1β, are pleiotropic cytokines with the host defense inflammatory responses [5,6]. Also, the induction and regulation of inflammatory responses has been shown to be under the control of key pro-inflammatory cytokines, including TNF-α and IL-1β [7] which are produced predominantly by antigen presenting cells, including macrophages. After then, numerous inflammatory mediators should serve to amplify and regulate innate immunity [8]. In tissue invading parasitic infections in immunodeficient mice, for example, with a filaria *Brugia* sp., pro-inflammatory cytokines, such as TNF-α and IL-1, are known to produce locally in parasitized lymphatics [9]. Therefore, in the present experiment, the author detected cytokine expressions in splenic dendritic cells and sera in order to clarify cellular immune responses in sparganosis of mice.

Spargana were collected from subcutaneous tissues of naturally infected snakes (*Rhabdophis tigrina*). After washing with sterile physiologic saline, worms were stored at 4˚C until infection to BALB/c mice. Mice were divided into normal and experimental groups (n = 3 for each group), from day 3 to week 8 post-infection (PI). Each mouse was infected by a single sparganum. After sacrificing mice, sera were collected and stored at -70˚C until cytokine assays. The spleen cells were digested with collagenase/EDTA solution. A positive magnetic bead using the mouse dendritic cell isolation kit (Miltenyi Biotec, Bergisch Gladbach, Germany) was reacted, and splenic DC were collected by passing over a Midi column (Miltenyi Biotec). RNA was prepared using RNA easy columns (Intron, Seoul, Korea). Primer sets of TNF-α and IL-1β were purchased from Bioneer (Cheongwon, Korea). Their primer information is listed in Table 1. All procedures were performed according to the manufacturer’s instructions. Briefly, RT-PCR reactions using the 1-step RT premix (Intron) were performed with an annealing temperature of 55˚C for 35 cycles. Expression of cytokines was cal-
culated with reference to the expression of β-actin and determined in duplicate. Serum cytokine levels were assayed using the mouse IL-1β EIA kit (IBL, Gunma, Japan), mouse TNF-α ELISA kit (Diaclone Research, Besancon, France) and measured duplicate. Results were expressed as a mean (ng/ml) ± SD concentration. Statistical significance was analyzed by the Student’s t-test (P < 0.05).

As shown in Fig. 1, the level of TNF-α increased rapidly and significantly and peaked at week 1 PI and that of IL-1β peaked at week 3 PI. However, TNF-α and IL-1β expressions showed a decreasing tendency at week 2 PI. In addition, serum TNF-α did not increase after infection and serum IL-1β increased slightly at week 2 PI and peaked at week 6 PI (Fig. 2). In this experiment, IL-1β appeared more important than TNF-α in host immune responses against mouse sparganosis.

Table 1. Primer sequences and amplicon size for TNF-α and IL-1β used in this study

| Name  | Amplicon size (bp) | Primer sequence |
|-------|-------------------|-----------------|
| β-actin | 395 | Forward 5'-AGG CTG TGC TGT CCC TGT ATG C-3' |
|       |       | Reverse 5'-ACC CAA GAA GGA AGG CTG GAA A-3' |
| IL-1β | 291  | Forward 5'-GCT ACC TGT GTC TTT CCC GTG G-3' |
|       |       | Reverse 5'-TTG TCG TTG CTT GGT TCT TG-3' |
| TNF-α | 300  | Forward 5'-GGG AGG TCT CTT TGA GTC CTA TG-3' |
|       |       | Reverse 5'-ACA TCC GAG GCT CCA GTG AAT TCG G-3' |

This selective and differential expression of pro-inflammatory cytokines may contribute to the host immune responses for attraction of effector cells to nearby sites of the parasites with
the capacity to limit the migration of tissue invading parasites, including spargana. On the contrary, the TNF-α response was preceded by the IL-1β response in the present study and this means that TNF-α may send signals to provoke IL-1β production in late phases of infection from immune cells. It is certain that the sparganum promote pro-inflammatory cytokine production and splenic DC innate immune responses are an essential first-line defense against the invading sparganum. In this study, serum TNF-α and IL-1β were reduced (P < 0.05) in early phases of infection when compared with normal control mouse sera. Such diminished production of IL-1β of immune cells is difficult to explain. Several reports [12,13] suggested that helminth parasite and their secreted products could modulate and suppress DC functions. An explanation have been issued by the presence of viable E. multilocularis metacestodes and possibly mediated by parasite’s excretory-secretory molecules that depressed the cytokine release and finally these molecules may prevent inflammatory reactions [11].

In conclusion, the current study provided evidence that the types and magnitudes of TNF-α and IL-1β responses in splenic DC and serum are chronologically regulated during the course of mouse sparganosis. It is suggested that certain pro-inflammatory cytokines may play an important role in the pathogenesis of mouse sparganosis.

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