NOTE  Anatomy

Change of calbindin D-28k protein expression in the mice hippocampus after lipopolysaccharide treatment

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ABSTRACT. A previous study showed that 1 mg/kg lipopolysaccharide (LPS) treatment did not lead to any neuronal death/degeneration (Received 1 October 2014/Accepted 5 November 2014/Published online in J-STAGE 26 November 2014)

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ABSTRACT. A previous study showed that 1 mg/kg lipopolysaccharide (LPS) treatment did not lead to any neuronal death/degeneration in the mouse hippocampus. In the present study, we examined the time-dependent changes of calbindin D-28k (CB) protein expression in the mouse hippocampus after a systemic administration of 1 mg/kg LPS. CB immunoreactivity was markedly increased in pyramidal cells of the hippocampal CA1/2 regions and in granule cells of the dentate gyrus from 3 hr to 48 hr after LPS treatment. At this point in time, CB protein level was also significantly increased in the mouse hippocampus. Thereafter, CB protein expression was decreased at 96 hr after LPS treatment. These results indicate that changes of CB protein expression may be associated with no neuronal death in the model of neuroinflammation with systemic administration of 1 mg/kg LPS.

KEYWORDS: calbindin, hippocampus, lipopolysaccharide

Calcium binding proteins, which act as calcium buffer, control the intracellular calcium homeostasis and cellular activity [3, 9]. Among the calcium binding proteins, calbindin D-28k (CB) binds to calcium ion with a fast association rate [4, 18]. In the brain, neuronal CB is thought to regulate intracellular response against various stimuli and provide neuroprotection against calcium-mediated neurotoxicity [2, 17]. In addition, CB-containing neurons play roles in memory, learning and long-term potentiation in the hippocampus [15].

Lipopolysaccharide (LPS), an endotoxin from gram-negative bacteria, leads to a strong response from normal animal immune system [16, 24]. LPS has been commonly used as a reagent for a model of systemic inflammatory response induced by infections, and systemic LPS treatment leads to neuroinflammation with neuroanatomical and neurochemical changes in some regions of the brain [8, 13]. Among the brain regions, the hippocampus is one of the brain regions most sensitive to LPS [6, 7].

Many previous studies have focused on effects of LPS on neuronal degeneration and immune response in various regions of the brain [19–21]. In addition, our previous studies showed that systemic administration of 1 mg/kg LPS led to microglial activation without any neuronal damage as well as changes of DNA repair ability in the mouse hippocampus [6, 11]. However, there is no study on LPS-induced changes of CB protein expression in the hippocampus. Therefore, in the present study, we examined the time-dependent changes of CB immunoreactivity and protein level in the mouse hippocampus after a systemic administration of 1 mg/kg LPS.

Six-week-old male ICR mice were purchased from Raon-Bio Inc. (Yongin, South Korea). The animals were housed in a conventional state under adequate temperature (23 ± 3°C) and relative humidity (55 ± 5%) control with a 12-hr light/dark cycle and provided with free access to food and water. The procedures for animal handling and care adhered to guidelines that are in compliance with the current international laws and policies (Guide for the Care and Use of Laboratory Animals, The National Academies Press, 8th Ed., 2011) and were approved by the Institutional Animal Care and Use Committee at Dankook University. All of the experiments were conducted to minimize the number of animals used and the suffering caused by the procedures used in the present study.

LPS (Sigma, St. Louis, MO, U.S.A.) was dissolved in saline and administered intraperitoneally with 1.0 mg/kg/10 ml/dose. The control animals were injected with the same volume of saline.

Mice (n=5 at each time point) were sacrificed at designated times (3, 6, 12, 24, 48 and 96 hr after LPS treatment). For the histological analysis, animals were anesthetized with zoletil 50 (30 mg/kg, Virbac, Carros, France) and perfused transcardially with 0.1 M phosphate-buffered saline (PBS, pH 7.4) followed by 4% paraformaldehyde in 0.1 M phosphate-buffer (PB, pH 7.4). The brain tissues were removed, cryoprotected and serially sectioned on a cryostat (Leica, Wetzlar, Germany) into 30 µm coronal sections, and they were then collected into 6-well plates containing PBS.

According to the method of our previous study [6, 11], immunohistochemical staining for CB was performed using rabbit anti-CB (1:200, Millipore, Temecula, CA, U.S.A.), biotinylated goat anti-rabbit IgG (1:200, Vector, Burlin-
game, CA, U.S.A.) and streptavidin peroxidase complex (1:200, Vector). In order to establish the specificity of the immunostaining, a negative control test was carried out with pre-immune serum instead of primary antibody. The negative control resulted in the absence of immunoreactivity in any structures (data not shown).

Six sections with 120 µm intervals per animal were selected to quantitatively analyze CB immunoreactivity. Digital images of the hippocampal subregions were captured with an AxioM2 light microscope (Carl Zeiss, Göttingen, Germany) equipped with a digital camera (Axiocam, Carl Zeiss) connected to a PC monitor. Semi-quantification of the immunostaining intensities in the pyramidal cells of the hippocampal CA1/2 region and in the granular cells of the dentate gyrus was evaluated with Image J 1.46 (National Institutes of Health, Bethesda, MD, U.S.A.). The mean intensity of immunostaining in each immunoreactive structure was measured by a 0–255 gray scale system (white to dark signal corresponded from 255 to 0). Based on this approach, the level of immunoreactivity was scaled as −, ±, + or ++, representing no staining, weakly positive, moderate or strong, respectively.

To confirm change in CB levels in the hippocampus at designated times (3, 48 and 96 hr after LPS treatment), animals at each time point (n=5) were used for western blot analysis according to the method of our previous study [6, 11]. In brief, after the hippocampus was homogenized and centrifuged, the supernatants were subjected to western blot analysis according to the method of our previous study [6, 11]. Therefore, it has been thought that neuronal death after systemic LPS treatment is dependent on the dose of LPS. Previously, we reported that 1 mg/kg LPS treatment induced cognitive impairment in mice [22, 23] and led to neuroinflammation, sickness and mild depressive-like behavior [5]. It has been also known that systemic LPS-induced neuronal death [19]. In addition, systemic LPS treatment induced cognitive impairment in mice [22, 23] and led to neuroinflammation, sickness and mild depressive-like behavior [5]. It has been also known that systemic inflammation induced by 10 mg/kg LPS treatment led to cellular apoptosis (15% neurons and 85% glia) in the rat hippocampus [21]. Previously, we reported that 1 mg/kg LPS treatment did not lead to the neuronal death/degeneration in the pyramidal and granule cells of the mouse hippocampus [6], although DNA repair ability in the hippocampal pyramidal and granule cells was affected after 1 mg/kg LPS treatment [11]. Therefore, it has been thought that neuronal death after systemic LPS treatment is dependent on the dose of LPS.

Table 1. Semi-quantifications of CB immunoreactivity in the mouse hippocampus after LPS treatment

| Groups | Control | LPS treatment |
|--------|---------|---------------|
|        | 3 hr    | 6 hr          | 12 hr | 24 hr | 48 hr | 96 hr |
| Pyramidal cells of CA1/2 region | ± | ++ | ++ | ++ | ++ | ± |
| Granule cells of dentate gyrus | + | ++ | ++ | ++ | ++ | ± |

Immunoreactivity is scaled as −, ±, + or ++, representing no staining, weakly positive, moderate or strong, respectively.
In the present study, we examined the changes of CB protein expression in the hippocampal CA1-3 regions and dentate gyrus after systemic treatment with 1 mg/kg LPS. It was reported that CB immunoreactivity was found predominantly in the pyramidal cells of the CA1/2 regions, not CA3 region, and in the granule cells of the dentate gyrus of the rat hippocampus [14]. We observed that the CB immunoreactivity in the pyramidal cells of the CA1/2 regions and in the granule cells of the dentate gyrus as well as the CB protein level in the hippocampus was markedly increased from 3 hr to 48 hr after LPS treatment. Although it is hard to exactly explain why CB protein expression was increased in the hippocampus after LPS treatment, it can be postulated that the increase of CB immunoreactivity in pyramidal and granule cells may be related to neuroprotection against LPS-induced neuronal death/degeneration in the mouse hippocampus, based on the results of the previous study [19]. In addition, we observed that CB immunoreactivity was increased in the stratum lucidum of the hippocampal CA3 region. It has been known that CB is closely associated with long-term potentiation, synaptic plasticity and memory functions [1, 10, 14]. Therefore, this result indicates that increase of CB immunoreactivity in the stratum lucidum may be related to the LPS-induced changes of hippocampal function, although we did not examine the changes of long-term potentiation, synaptic plasticity and memory functions after LPS treatment in the present study.

In conclusion, CB protein expression was apparently increased in the mouse hippocampus after systemic administration of 1 mg/kg LPS. These results indicate that changes in CB protein expression may be associated with no neuronal death in the model of neuroinflammation with systemic administration of 1 mg/kg LPS.
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