Diurnal Spatial Rearrangement of Microglial Processes through the Rhythmic Expression of P2Y_{12} Receptors

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Abstract

Microglia plays important roles in synaptic reorganization during the postnatal developmental stage. Moreover, microglia continuously surveys the functional state of the synapse and change to improve the function. This phenomenon was attributed to the fine process of extension and retraction. However, the mechanism underlying the dynamics of microglial movement and function is still unclear. We herein report that cortical microglia exhibit clock gene-regulated diurnal morphological changes. Cortical microglia extended their processes during the dark phase and retracted them during the light phase. These diurnal changes were also observed in cortical microglia from animals housed under constant darkness, but not in cortical microglia from clock-mutant mice. The mean contact ratio of the microglia-synapse interactions was significantly larger during the dark phase than the light phase. These diurnal changes in microglial morphology and microglia-synaptic interactions were significantly inhibited by the systemic administration of clopidogrel, a P2Y_{12} receptor (P2Y_{12}R) blocker. We further observed diurnal variation in the P2Y_{12}R expression in cortical microglia. The report analyses further revealed that P2Y_{12}R was regulated by a negative feedback loop of the clock system. These observations suggest that the microglial clock system drives the diurnal morphological changes of microglia and microglia-synapse interactions by controlling the P2Y_{12}R expression.

Keywords: Microglia; Microglia-synapse interaction; P2Y_{12} receptor; Clopidogrel; Clock genes

Introduction

Several lines of evidence suggest that microglia continuously move their processes by extension and retraction [1,2]. The motile fine processes of microglia are directly contact with synaptic structures [3], implying that microglia regulate the functional state of neurotransmission in the normal brain. During the postnatal developmental stage, microglia have an important role in the regulation of synapse maturation with phagocytosis occurring through CX3CR1 signaling in the hippocampus or barrel cortex [4,5]. In addition, microglia engulfs and eliminates unnecessary synapses in a complement-dependent manner in the dorsal lateral geniculate nucleus [6]. Microglia also eliminates dendritic spines to adapt to environmental alterations in the visual cortex [7]. These observations strongly suggest that microglial processes have important roles in the fine tuning of the neural circuit in the healthy brain.

Microglial processes contact synaptic sites by their recognition of neuronal activity. A reduction of the neuronal activity in the visual cortex after an injection of tetrodotoxin into the retina resulted in a decreased frequency of contacts between microglial processes and synaptic sites [3]. On the other hand, the neuronal activity is continuously changing during the day. It is well known that the suprachiasmatic nucleus exhibits a daily oscillation in neuronal activity, which regulates the circadian rhythm [8]. Diurnal neuronal activity was also reported in the cerebral cortex [9,10]. Furthermore, the total net change in the dendritic spine is also regulated by the sleep-wake cycle [11,12]. These observations prompted us to examine the possible involvement of microglia in the diurnal changes in the neuronal synaptic activity through microglia-synapse interactions. Therefore, in the present study, we examined the possible association of the diurnal changes with the microglia-synapse interactions using a quantitative morphological approach. We herein provide the first evidence of the diurnal morphological changes in microglia and microglia-synapse interactions, which are regulated by clock gene-driven expression of P2Y_{12} receptors (P2Y_{12}R).

Materials and Methods

Animals

All efforts were made to minimize animal suffering and to reduce the number of animals used. Eight-week-old male C57BL/6 mice and Clock-mutant (Clock/Clock) mice, which have a point mutation in exon 19 of the Clock gene (Jackson Laboratory, Bar Harbor, ME, USA), on an ICR background were purchased from Charles River Japan, Inc. (Kanagawa, Japan). The mice were housed under either normal light/dark (LD) condition (light on at 07:00 and light off at 19:00) or constant darkness (DD) in a temperature-controlled room (24 ± 1°C) with a humidity of 60 ± 10%, and were provided food and water ad libitum. They were adapted to the LD for two weeks before the experiments. Under the LD condition, the ZT0 (Zeitgeber time) was designated as lights on and ZT12 as lights off. For mice housed under the DD condition for one day, the circadian time (CT) was used instead of ZT, and CT0 was characterized as the beginning of the subjective light phase, and CT12 was defined as the beginning of the subjective dark phase. All the preparations were performed under the red light. All animals were treated in accordance with the guidelines stipulated by the animal care and use committee of Kyushu University.

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Isolation of microglia from adult animals

Wild-type and clock-mutant mice were anesthetized and perfused transcardially with phosphate buffered saline (PBS), then the mice were decapitated at the time point of ZT2, 6, 10, 14, 18 or 22 or at CT2 or14 (n=3 at each time point). Brain tissue samples were cut into small pieces and single-cell suspensions were obtained by enzymatic digestion using the Neural Tissue Dissociation Kit (Papain). The tissue was further mechanically dissociated using a wide-tipped, fire-polished Pasteur pipette and the suspension was applied to a 30 µm cell strainer. The cells were magnetically labeled with CD11b MicroBeads. The cell suspension was loaded onto a magnetic cell sorting (MACS) column placed in the magnetic field of a MACS Separator, and the negative fraction was collected. After removing the magnetic field, the CD11b-positive cells were eluted as a positive fraction. Isolated cells were incubated with anti-mouse CD45-fluorescein (FITC) and anti-mouse CD11b-phycocerythrin (PE) at optimal dilutions. Stained cells were analyzed by a fluorescent activated cell sorting (FACS) (Becton-Dickinson) using the Cell Quest software program. The purity of microglia was calculated to be 78.24%. All reagents were purchased from Miltenyi Biotec. All manipulations were terminated within 3 h in the present study to prevent any effect of microglial activation due to the isolation procedure.

Quantitative RT-PCR analysis

Total RNA was extracted from microglia cells using RNAiso (Takara Bio Inc., Shiga, Japan) at ZT2, 6, 10, 14, 18 or 22 or at CT2 or 14. Complementary DNA (cDNA) was prepared via reverse transcription of the total RNA using a ReverTra Ace qPCR RT kit (Toyobo Co. Ltd., Osaka, Japan). Diluted cDNA samples were analyzed by real-time RT-PCR. Real-time PCR was performed using the THUNDERBIRD™ SYBR® qPCR Mix (Toyobo) and the 7500 Real-time PCR system (Applied Biosystems, Foster City, CA). To determine the copy numbers, we chose the 2^-ΔΔCt method using a calibrator as described previously [13]. The primer sequences used for the quantitative PCR of target gene (P2Y12R) were forward: 5’-CACAGAGGCCTTTGGGAACCTA-3’ and reverse: 5’-TGCTCTCCTGCTCTGTAAC-3’ and for β-actin (as a control) were forward: 5’-CACACCTCTTACATAAGCTGTCG-3’ and reverse: 5’-CATGATCTGCTGATCTTCTCA-3’ and the resulting amplicons were 126 bp and 109bp, respectively. These primers were designed to flank a region that contains at least one intron.

Construction of reporter and expression vectors

To contrast the luciferase reporter vector for the P2Y12R gene (P2Y12R-Luc), a 5’-upstream fragment of the promoter region of the mouse P2Y12R gene spanning bp -944 to +51 (the number is the distance from the putative transcription starting site, +1; Genbank Accession #NM_027571) was amplified by PCR, and the product was fused to the luciferase gene using a pGL3-promoter vector (Promega). The expression plasmids for clock and clock-related genes were prepared as follows: the coding regions of clock or clock-related genes were obtained by RT-PCR and used after their sequences were confirmed, as reported previously [14]. All coding regions were ligated into the pcDNA3.1 vector (Invitrogen).

Luciferase reporter assay

The NIH3T3 cells were maintained in DMEM supplemented with 5% FBS at 37°C in a humidified 5% CO2 atmosphere [15]. The NIH3T3 cells do not express the P2Y12R gene in NIH3T3 cells (data not shown). The day before transfection, cells were seeded (1×10^5 cells/well) into 24 well plates. The next day, the cells were transfected with 100 ng of the reporter construct and 0.5 µg (total) of expression constructs using the Lipofectamine-LTX reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's instructions. To correct for the variations in transfection efficiency, 0.5 ng of pGL4.74 [hRluc/TK] vector (Promega, San Luis Obispo, CA, USA) was co-transfected into each well. The total amount of DNA per well was adjusted by adding pcDNA3.1 vector (Invitrogen). At 48 h post-transfection, the cells were harvested with 200 µL passive lysis buffer, and 50 µL of the extracts were used for firefly luciferase and Renilla luciferase assays by luminometry. The ratio of firefly luciferase activity expressed from reporter plasmids to the Renilla luciferase activity expressed from pGL4.74 [hRluc/TK] in each sample served as a measure of the normalized luciferase activity, as reported previously [14].

Histology

Mouse tissues were examined by an immunohistochemical analysis as described previously [16]. All mice were deeply anesthetized with sodium pentobarbital (50 mg/kg i.p.) and perfused transcardially with 0.1 M phosphate buffer (PB; pH 7.4), followed by 4% paraformaldehyde (PFA) in 0.1 M PB at ZT2, ZT14, CT2 and CT14 (n=3 animals at each time point) for immunohistochemistry. Transverse cortical sections (free floating, 50 µm thick) were prepared using a vibratome (VT1000S, Leica, Nussloch, Germany). The floating sections were incubated with a mixture of 1.0% BSA in PBS containing 0.05% sodium azide 1 h at room temperature, then the sections were stained with an antibody against Iba1 (1:10,000; Wako) for 7 d at 4°C and with Cy3 conjugated donkey anti-rabbit IgG (1:400, Jackson Immunoresearch) for 3 h at 4°C. The sections were then washed with PBS and mounted in the Vectashield anti-fading medium (Vector laboratories, Peterborough, UK) and examined with a confocal laser-scanning microscope (CLSM, LSM510MET, Carl Zeiss, Jena, Germany). The images of microglia were captured from layer 2/3 in the primary somatosensory cortex.

Intracellular labeling and subsequent Iba1 staining

The cortical sections perfused with 4% PFA were sectioned into 100 µm thick samples by a vibratome (VT1000S, Leica, Nussloch, Germany) (n=24-26 slices from three animals). The sections were incubated with YOYO-1 (1:10,000; Molecular Probes) for 3 min to visualize the cell body. A glass electrode filled with 4% lucifer yellow was applied to the YOYO-1-positive cell bodies under a microscope equipped with a 40×water immersion objective by using a laser with a wavelength of 488 nm. When the dendrite apparently changed brightness following the insertion of the glass electrode into the cell body, current was injected (up to -10 nA) in the current clamp mode to further inject the lucifer yellow, as was previously described for a similar protocol [17]. After the injection of the lucifer yellow, the sections were incubated with 4% PFA for 24 h at 4°C. Next, the sections were further stained with an anti-Iba1 antibody as described above. The images were captured by a CLSM at wavelengths of 477 nm and 543 nm.

Quantitative morphological analyses of microglia and dendritic spines

All images were processed using the Image J software program (National Institutes of Health, Bethesda, MD, USA). The acquisition of 3D images was carried out by CLSM with 63×objective fields for single staining with Iba1 and with 100×objective fields for counter-staining with Iba1. Stacks of 70-90 serial optical sections were obtained from...
each cerebral cortex (22-27 dendrites from 10-15 cells from three animals). The morphological analyses of microglia were performed on Z-projections of confocal images. The cell number of microglia was counted for Z-projection images with a thickness of 50 µm within 0.01 mm². Microglial processes were traced and then reconstructed as single microglial images using the Simple Neurite Tracer software program plug-in. Semi-automated tracing of dendrites was performed for the 3D image data [18], which were used to calculate the total length of microglial processes and the branch number. The topological skeletonized images of single microglia were converted from a binarized image of single microglia using the skeletonize function. The spine density was determined by manually identifying the spines and using 3D images to measure the dendrite length with the Simple Neurite Tracer plug-in. The number of dendritic spines in close apposition with microglial processes or enwrapped by microglial processes was counted. The number of dendritic spines contacted with microglial processes was normalized to the total number of dendritic spines counted and represented as the contact ratio. 18-20 microglial cells were analyzed in each animal. The 3D illustrations of microglia and spines were made using the Reconstruct 1.1 software program (http://synapses.clm.utexas.edu).

Immunoblotting

The samples were separated by electrophoresis on 10% SDS-polyacrylamide gels (n=3 independent experiments). Proteins on SDS gels were transferred electrophoretically to nitrocellulose membranes. The membranes were washed with PBS and incubated at 4°C overnight under gentle agitation with each primary antibody: anti-P2Y₁₂R (1:1,000; Novus Biologicals) and anti-β-actin (1:1,000; Santa Cruz). After washing, the membranes were incubated with horseradish peroxidase (HRP)-labeled anti-goat (1:1,000; R&D Systems) or anti-rabbit (1:1,000; GE Healthcare) antibodies for 1 h at room temperature. Subsequently, the membrane-bound, HRP-labeled antibodies were detected using an enhanced chemiluminescence detection system (ECK lit; GE Healthcare) with an image analyzer (LAS-4010; Fuji Photo Film).

Drug administration

Clopidogrel (25 mg/kg, LKT Laboratories, St. Paul, MN), a selective P2Y₁₂R inhibitor, was systematically administered three times every 3 h from ZT10-18 and CT10-18. Then, specimens for the histological experiments were prepared at ZT2 and CT2. In some experiments, the mice were decapitated at ZT14 and CT14 after either saline or clopidogrel administration at ZT10 and CT10.

Statistical analyses

All data are shown as the means ± SEM. The statistical analyses were performed using two-tailed unpaired Student’s t-tests, a one-way analysis of variance (ANOVA) with a post-hoc Tukey’s test or a two-way ANOVA with repeated measurements using the GraphPad Prism Software package (GraphPad Software Inc., San Diego, CA, USA). Unless otherwise indicated, data met the assumptions of equal variances. Differences were considered to be significant at p<0.05.

Results

Diurnal morphological change in cortical microglia

The quantitative morphological analyses of cortical microglia in the young adult C57BL/6 mice were performed using the skeletonized images reconstructed from Z-stack images of Iba1-immunostained microglia (Figures 1A-1D). Diurnal variation in spatial area and structural complexity were observed in cortical microglia (Figures 1E-1H). The mean total length of processes (Figure 1I) and the mean number of branch points (Figure 1J) of cortical microglia

![Figure 1](image-url)
Regulation of microglia-synapse interactions by P2Y12R

P2Y12 receptors (P2Y12R), which are G\text/sub>i/o-coupled ATP receptors expressed exclusively on microglia in the brain [20], regulate microglial process extension [21]. The loss of P2Y12R expression accompanies microglial transformation from a highly ramified to amoeboid state following neuronal injury [21]. Therefore, the possible involvement of P2Y12R in the diurnal variation of the microglial morphological complexity was next examined using clopidogrel, a P2Y12R blocker that can cross the blood-brain barrier after systemic administration [22]. The extension of microglial processes during the dark phase (ZT14) was significantly inhibited by the systemic treatment with clopidogrel (Figures 3A-3C). Lucifer yellow, combined with subsequent Iba1-staining showed that the microglial processes frequently made contact with dendritic spines (Figure 3D). The three-dimensional (3D) reconstructed images showed that microglial processes wrapped (Figures 3E-3H) and apposed (Figures 3I-3K) the dendritic spines. The mean contact ratio of microglia and dendritic spines was significantly higher at ZT14 (4.72 ± 0.35%) than at ZT2 (3.07 ± 0.54%) (Figure 3I), and correlated with the microglial complexity (Figures 1I and 1J). The method used in this study may underestimate the contact ratio of microglia and dendritic spines, because microglial processes are highly motile and make only a brief contact with dendritic spines [3]. Furthermore, the increase in dendritic spine density at ZT14 (0.85 ± 0.03/µm) compared with ZT2 (0.66 ± 0.03/µm) was similar to the previously reported [11]. More interestingly, systemic administration of clopidogrel suppressed the contact ratio at ZT14 (Figure 3I; 3.05 ± 0.33%) and preserved the dark phase-like dendritic spine density even at ZT2 (0.84 ± 0.02/µm). These observations suggest that P2Y12R plays a key role in microglial interactions with synaptic structures.

Clock gene-regulated expression of P2Y12R in cortical microglia

To examine whether cortical microglia exhibit clock gene expression, which would regulate P2Y12R gene expression, microglia were isolated from the adult mouse cerebral cortex by a magnetic cell sorting method using CD11b microbeads at 4 h-intervals over a 24 h period. Microglia was found to exhibit a circadian oscillation of P2Y12R transcripts with a peak at ZT14 (Figures 4A and 4B). Furthermore, a rhythmic pattern of P2Y12R was also observed in microglia isolated from clock gene-regulated cortical microglia, we utilized clock-mutant (Clock/Clock) mice, which exhibit disrupted circadian gene expression [19]. Microglia in the somatosensory cortex of wild-type ICR mice also exhibited the diurnal morphological change of microglia (Figures 2A, 2B, 2E, 2F, 2I and 2J). In contrast, there was no diurnal morphological change of cortical microglia in clock-mutant mice (Figure 2C, 2D, 2G, 2H-2J).

Figure 2: The deficits in the circadian variation of the spatial area and the structural complexity of cortical microglia from clock-mutant mice. A-D: CLSM stack images of cortical microglia at ZT2 (A) and ZT14 (B) from wild-type mice and ZT2 (C) and ZT14 (D) from clock-mutant mice. E-H: The topological skeletonized reconstructed images from the 2D images of cortical microglia at ZT2 (E) and ZT14 (F) from wild-type mice and ZT2 (G) and ZT14 (H) from clock-mutant mice. Scale bars, 10 µm. I, J: The clock mutation led to a total loss of the alterations in the process length (I) and branch points (J) of cortical microglia. The data are the means ± S.E.M. (animals=3, n=27-29 sections each).

The asterisks indicate statistically significant differences between values (*, P<0.05; **, P<0.01; one-way ANOVA). K: The number of cortical microglia within a 0.01 mm² area was not significantly different at each time point (animals=3, n=27-29 sections each).
under the DD conditions (Figure 4C). In addition, the diurnal variation in the expression of P2Y$_{12}$R occurred in a clock-dependent manner (Figures 4B and 4C). The protein level of P2Y$_{12}$R in cortical microglia was also increased at ZT14 compared to ZT2 (Figure 4D).

The E-box site (CACGTG), the target of the circadian clock effectors BMAL1 and CLOCK, is located 382 bp upstream of P2Y$_{12}$R. Therefore, the relationship between the clock genes and P2Y$_{12}$R was further analyzed using a luciferase reporter assay system targeting this region (Figure 4E). CLOCK/BMAL1 proteins enhanced the transcriptional activity of P2Y$_{12}$R-Luc that contained the E-box, whereas CLOCKΔ19/BMAL1 did not (Figure 4F). In addition, the CLOCK/BMAL1-mediated transactivation of P2Y$_{12}$R-Luc was significantly repressed by the addition of PER2 or CRY1 (Figure 4G). These observations indicate that the diurnal changes in the P2Y$_{12}$R gene expression are regulated by the molecular clock system, consisting of a negative feedback loop with the CLOCK/BMAL1 complex binding to E-boxes in its promoter region.

**Discussion**

The major finding of this study is that cortical microglia exhibit diurnal morphological changes, which are regulated by the clock gene-driven diurnal expression of P2Y$_{12}$R. During the dark phase, microglia extends their processes resulting from an increased level of P2Y$_{12}$R expression. During the light phase, the decrease in the P2Y$_{12}$R level results in the partial retraction of microglial processes. Furthermore, this diurnal change in the microglial morphology influences the ratio of microglia-synapse interactions. It is though those microglia extend their processes specifically toward highly active synapses that release ATP and glutamate, because a reduction in the neuronal activity leads to the suppression of contact between microglial processes and dendritic spines [3]. Furthermore, sensory deprivation started the retraction of microglial process over 6 hours. Both ATP and glutamate increase the morphological complexity in microglia, such as the extension of their process and elevation of the number of branch points [23]. Glutamate itself also induces the release of ATP from microglia [23] and astrocytes [24]. ATP induces the ramification [25] and sprouting of new processes from existing microglial processes [26]. Consistent with these observations, apyrase, an ATP hydrolyzing enzyme, decreases the total number of branch points [23]. In our study, clopidogrel, a P2Y$_{12}$R inhibitor, suppressed the circadian microglial process extension. Besides the circadian oscillation of P2Y$_{12}$R gene expression, extracellular ATP concentration secreted from neurons and astrocytes was also known to exhibit circadian oscillation [27,28]. Therefore, the circadian oscillation of the extracellular ATP concentration should be also considered as a causative factor for diurnal morphological variation of cortical microglia. These observations strongly suggest that ATP regulates both the length and structural complexity of microglial processes through P2Y$_{12}$R. Further studies are still needed to clarify the involvement of endogenous ATP on the process extension of microglia, and would be facilitated by the development of a blood-brain barrier permeable ATP hydrolyzing enzyme.

Although clopidogrel is used clinically as an anti-platelet agent, clopidogrel or its metabolite(s) can cross the blood-brain barrier [29]. Furthermore, effects of clopidogrel on spinal microglia were previously observed after systemic administration [22]. The expression
of P2Y12R is limited to microglia, and the receptors are not expressed on neurons or astrocytes [20,21,30]. Therefore, it is considered that the systemic administration of clopidogrel can specifically block the P2Y12R expressed on microglia. The concentration of clopidogrel used in our study is extremely high compared to the clinical dose. The usual clinical dose of clopidogrel is one 75 mg tablet per day. This concentration is equivalent to approximately 1 mg/kg in an adult human with a body weight of 60 kg. In the present study, we administrated 25 mg/kg of clopidogrel three times a day. Therefore, it is unlikely that the usual clinical dose of clopidogrel would inhibit the microglia-synapse interactions through a blockade of the P2Y12R expressed on microglia.

During the phagocytic elimination of dead cells in the dentate gyrus, microglia formed a ball-and-chain structure on their terminal processes [31]. Furthermore, the synaptic structure is phagocytosed by microglial processes during the postnatal developmental stage [4,6] and is involved in the adaptation to new environmental situations [7]. In this study, we found that the mean density of dendritic spines of the cortical neurons was significantly larger during the dark phase (ZT2) than the light phase (ZT14). Furthermore, the mean density of dendritic spines during the light phase was significantly increased by microglial processes during the postnatal developmental stage [4,6].

Conclusion
We have herein provided evidence of a novel system wherein the microglial process dynamics are regulated by the intrinsic clock system. The intrinsic microglial molecular clock drives the expression of P2Y12R during the dark phase to extend their processes and increase contacts with dendritic spines, which may, in turn, decrease the density of dendritic spines for the subsequent light phase. It has been proposed that dysfunction of the clock system is one of the risk factors for the psychiatric diseases. In addition, abnormal microglia has a significant association with neurological disorders [33,34]. Therefore, the microglial clock may provide valuable targets for the development of novel therapeutic agent for the neurological disorders, and further research on this topic will aid in understanding the function and dysfunction of the brain.

Disclosure
The author reports no conflicts of interest associated with this work.

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