Review Article

The Role of Mammalian Glial Cells in Circadian Rhythm Regulation

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Received 14 September 2017; Accepted 19 November 2017; Published 25 December 2017

Circadian rhythms are biological oscillations with a period of about 24 hours. These rhythms are maintained by an innate genetically determined time-keeping system called the circadian clock. A large number of the proteins involved in the regulation of this clock are transcription factors controlling rhythmic transcription of so-called clock-controlled genes, which participate in a plethora of physiological functions in the organism. In the brain, several areas, besides the suprachiasmatic nucleus, harbor functional clocks characterized by a well-defined time pattern of clock gene expression. This expression rhythm is not restricted to neurons but is also present in glia, suggesting that these cells are involved in circadian rhythmicity. However, only certain glial cells fulfill the criteria to be called glial clocks, namely, to display molecular oscillators based on the canonical clock protein PERIOD, which depends on the suprachiasmatic nucleus for their synchronization. In this contribution, we summarize the current information about activity of the clock genes in glial cells, their potential role as oscillators as well as clinical implications.

1. Introduction

Most light-sensitive organisms have built-in time-measuring devices that are commonly known as circadian clocks. The term \textit{circadian} was introduced by Halberg to describe the biological rhythms that have a period of approximately 24 h and are known as circadian rhythms [1]. Circadian rhythms are present along the phylogenetic scale, in mammals regulate a plethora of functions such as the rest-activity cycle, hormone secretion, and daily variations in metabolism and body temperature [2].

The intracellular circadian clock is based on a transcription-translation feedback system that drive the self-sustaining clock mechanism in the suprachiasmatic nucleus (SCN, the “master clock”) and in peripheral tissues (“peripheral clocks”) [3, 4]. At the molecular level, the molecular machinery that generates circadian rhythms involves CLOCK- (circadian locomotor output cycles kaput) BMAL1 (brain and muscle aryl hydrocarbon receptor nuclear translocator-like protein 1) heterodimers that control the periodic expression of Per (periods 1–3) and Cry (cryptochrome 1,2) genes. These gene products form the PER-CRY heterodimer that is translocated to the nucleus inhibiting their own transcription. Additionally, an accessory regulatory loop involves the rhythmic regulation of Bmal1 transcription through the coordinated action of the transcriptional repressor REV-ERBa (Reverse Erb alpha) and the transcriptional activator RORα (retinoid-related orphan receptor-alpha) [5–8].

In mammals, the SCN synchronizes multiple peripheral clocks, in numerous tissues and cell types, presumably via the combination of neural and humoral signaling [2, 9]. The general consensus of the cellular identity of the oscillating cells in the brain points to the neurons, although it has been demonstrated that the glial cells are circadian oscillators as well, and their synchronization also depends on the SCN [10–12].
Glial cells make up a large fraction of human nervous system cells, with numbers exceeding those of neurons by a factor of ten, depending on the brain structure studied. Particularly, glial cells occupy about half the volume of the brain and participate in diverse functions, including regulation of synaptic transmission, plasticity, behavior, and synapse development, and these cells are also involved in neurodegeneration [13–17]. Interestingly, it has been described that glial cells also play an important role in the regulation of circadian rhythms [18–23], although little attention has been paid to this function. Accordingly, we summarize here the recent findings about clock genes in glial cells, the plausible role of the glial cells as cellular oscillators, and the possible medical implications of clock genes in this cell type.

2. Clock Genes in Glial Cells

2.1. Astrocytes. This type of glial cell is involved in the buffering of extracellular K⁺, regulating neurotransmitter release [24], forming the blood-brain barrier, releasing growth factors, and the regulation of complex brain mechanisms, such as sleep homeostasis [25] and memory [26–28], among other functions.

In 1990, it was suggested that glial cells might express molecular oscillators, which are based on the clock protein PER. Particularly, it was demonstrated that PER was localized both in neurons and glial cells of the fly brain, which showed robust circadian rhythms and abundance [11]. Subsequently, Ewer and coworkers reported that certain weakly rhythmic flies contained detectable PER only in glia, suggesting that glial oscillators play an important role in the pacemaker driving rhythmic behavior [12]. Later, it was confirmed the rhythmic expression of clock genes in rat and mouse astrocytes, indicating that these cells contain a PER-based molecular oscillator that dampens in the absence of neuronal signals [29, 30]. These astroglial cultures were capable to display a sustained rhythmicity for 7 days when cocultured with SCN explants, whereas cortical explants did not influence rhythmicity [29], suggesting that the presence of sustained rhythms in glial cells requires a secreted neuronal factor expressed in the SCN. Temperature cycles entrain Per1 rhythms in astroglial cultures [29] however are unlikely to be a relevant factor, since exposure to SCN explants sustained glial rhythms without any change in temperature.

Several studies have explored the role of the mammalian PER-based oscillator in glial physiology. It has been reported that Glast (glutamate/aspartate transporter) expression and protein levels within the SCN present a diurnal rhythm in a light/dark (12/12 h) cycle [31]. However, it was not determined whether this rhythm persist or not in nonrhythmic conditions (constant darkness or constant light), in other words in lack of environmental information. Results of Spangagel and collaborators are complementary with the observation concerning to GLAST levels, which do not display an obvious rhythmicity in the Per2 mutant mice pointing out the presence of a circadian control [31]. Beaulé and colleagues reported that cultured cortical astrocytes from Clock mutant animals have reduced Glast mRNA and protein levels [32], proposing that the vast majority of glial glutamate (Glu) uptake activity is a function of the transcription factors Clock and Npas2 (neuronal PAS domain protein 2) and of the transcriptional regulator Per2 [32, 33]. This dependence could be explained by the involvement of CLOCK and NPAS2 in the indirect regulation of Glast transcription or in GLAST protein stabilization and/or localization [34]. It should be noted that no evidence has been demonstrated for circadian changes in Glu uptake, suggesting a noncircadian role for clock proteins that might be involved in the regulation of Glast gene transcription or Glast mRNA translation and/or stability [32, 33].

Concerning Glu, it is known that this neurotransmitter participates in photic entrainment of circadian rhythms. In 2015, it was reported that in cultured Bergmann glial cells, Bmal1 expression is Glu time- and dose-dependent. This phenomena might be a result of stabilization of the protein after it has been phosphorylated by PKA (cyclic AMP-dependent protein kinase) and/or PKC (Ca²⁺/diacylglycerol-dependent protein kinase), pointing out that Glu is critically involved in glial Bmal1 expression and that glial cells are important in the control of circadian rhythms in the cerebellum [22].

It has been recently demonstrated that not only SCN neurons but also SCN astrocytes possess pacemaking properties [35]. By using long-term live imaging, Brancaccio and colleagues simultaneously codetected circadian oscillations of neuronal and astrocytic intracellular calcium ([Ca²⁺]i) within the SCN, with [Ca²⁺]i levels peaking during the circadian day and night. Thereby, these oscillations of [Ca²⁺]i were antiphasic and showed a complementary waveform [35]. In the same study, it was reported widespread circadian oscillations of extracellular Glu ([Glu]e) in the SCN in phase with astrocytic [Ca²⁺]i. These circadian oscillations of [Glu]e are generated intrinsically in the SCN and also depend directly on astrocytic metabolism [35]. Using pharmacological inhibition of the glial and neuronal isoforms of the Glu transporters, a continuing circadian oscillation of Glu release by astrocytes is observed. Remarkably, [Glu]e oscillations are generated by concerted rhythms of release and uptake, and blocking Glu uptake impairs the fine-tuning of the [Glu]e/[Ca²⁺]i relationship, reducing the robustness of the rhythms of neuronal [Ca²⁺]i across the SCN. Consequently, SCN cellular oscillators progressively desynchronize, until the [Glu]e/[Ca²⁺]i alignment is restored. Presynaptic NMDA (N-methyl-D-aspartate) receptors 2C-mediated glutamatergic glutotransmission inhibit neuronal activity during circadian night, and this mechanism is essential to sustain circadian rhythmicity in the dorsal SCN [35]. Accordingly, during the circadian night, SCN astrocytes are metabolically active (high [Ca²⁺]i) and release high levels (baseline activity) of Glu into the extracellular space, which in turn activates presynaptic NR2C-expressing neurons in the dorsal SCN, thereby increasing GABAergic inhibitory tone across the circuit. In contrast, during the circadian day, [Glu]e is reduced by diminished glial release and increased EAAT-mediated Glu uptake and consequently, GABAergic tone is reduced, thereby derepressing spontaneous membrane potential, neuronal [Ca²⁺]i, and facilitating electrical firing [35].
Moreover, it was reported that SCN astrocytes are functional circadian oscillators, which modulate the period of SCN and the rest-activity rhythms [36]. The loss of rhythm in SCN astrocytes by Bmal1 deletion leads to an extended circadian period of rest-activity rhythms. This Bmal1 deletion in a small proportion of SCN cells appears to change the period of the SCN and behavior by the loss of rhythmicity in 20% of SCN cells that express AVP (arginine-vasopressin) or 10% of cells that express Aldh1l1 (specific astrocytic marker) or GFAP (glial fibrillary acid protein) [36].

Earlier studies in SCN astrocytes revealed high-amplitude daily rhythms in the expression of GFAP [37] and their coverage of the soma and dendrites of vasointestinal polypeptide- and AVP-expressing SCN neurons, which are related with modifications in synaptic innervation of these neurons [38, 39]. Rhythmic pattern of GFAP was observed in constant darkness in the SCN of hamsters, rats, and mice [37, 40], suggesting that these rhythms are intrinsic and independent of external light cues. Although the role of daily oscillations of GFAP in SCN is unknown, it has been associated with two main aspects of the clock functioning: metabolic exchanges and plasticity [38]. According to this last aspect, it has been demonstrated that mice lacking the Gfap show impaired long-term depression in the cerebellum, reduced eye-blink conditioning [41], longer periods of activity, and more arrhythmicity in constant light conditions compared to wild type [40, 42]. These results indicate that GFAP in glial cells plays a role in the regulation of neuronal function.

A daily variation of GFAP in the mouse SCN, as well as the NF-κB (nuclear factor-κB) expression in SCN astrocytes has been documented using tissue slices and primary cell cultures [43]. Particularly, in the latter case, LPS (lipopolysaccharide), IL-1α (interleukin-1 alpha), and TNFx (tumor necrosis factor alpha) promoted the activation of NF-κB, indicating that SCN astrocytes mediate the input signals to the circadian system from the immune system via NF-κB signaling [43].

ROαα is expressed in astrocytes but not in microglia. Studies using staggerer mice, which have a 122 bp deletion in the ROαα gene, allowed the identification of several functions of this nuclear receptor, both in the periphery and in the CNS. Interestingly, a massive cerebellar neurodegeneration leading to severe ataxia was also observed in these mice [44].

Furthermore, it has also been reported that ROαα-deficient mice have abnormal immune responses, associated with increased levels of IL-1β, IL-6, and TNFα [45]. An additional report showed that in primary astrocyte cultures, ROααs directly participate in the regulation of the inflammatory reaction via the inhibition of the NF-κB pathway. Thus, in a noninflammatory condition, the nuclear receptor directly increases IL-6 expression, while in an inflammatory condition, ROαα reduces cytokine-induced Il-6 upregulation [46].

Other nuclear receptors involved in the inflammatory response are REV-ERBα and REV-ERBβ; both receptors are expressed in rat C6 cells and in astrocyte cultures derived from rat cortex and spinal cord [47]. Particularly, in rat C6 astroglial cells, it has been reported that TNF significantly increases chemokine Ccl-2 (monocyte chemoattractant protein-1), IL-6, iNOS (inducible nitric oxide synthase), and Mmp-9 (matrix metalloprotease-9) mRNA levels. However, both isoforms of REV-ERB inhibit TNF-induced upregulation of Ccl-2 and Mmp-9 mRNA levels. Particularly, REV-ERBα and REV-ERBβ decrease MMP-9 expression via HDAC3 (histone deacetylase 3) [47]. Moreover, it has been shown that REV-ERBα inhibits Il-6 upregulation in murine skeletal muscle cells and macrophages [48–50], suggesting that the activity of this nuclear receptor is tissue-specific.

Circadian expression of clock genes such as Per1, Per2, Cry1, and Bmal1 can be observed in mice spinal cord. Surprisingly, circadian expression of Gs (glutamine synthetase, a glial-enriched enzyme) and COX-1 (cyclooxygenase-1) at both mRNA and protein levels was also detected in the same brain area [51]. Moreover, circadian changes in the expression of Gs suggest that astrocyte metabolism is subjected to circadian modulation. Whereas, the disruption of astroglial function using fluorocitrate (a glial metabolic inhibitor) led to the suppression of the oscillating expression of not only Gs and COX-1 but also the expression of clock genes. These findings suggest that spinal circadian expression of clock genes depends on the activity of astrocytes, since the inhibition of astrocytic function disrupts circadian gene mRNA expression [51].

Gliotransmission is the process by which astrocytes communicate with immediate glia and neurons through the release of transmitters such as ATP and Glu [42, 52, 53]. In vivo, a circadian pattern of ATP release appears to derive primarily from astrocytes within the SCN; however, the functional implications of these extracellular ATP rhythms are unknown [54]. Moreover, it has been shown that astrocytes display daily extracellular ATP oscillations that rely on key clock genes (Clock, Per1, and Per2) and inositol triphosphate signaling [55], suggesting that extracellular ATP levels are augmented at specific hours of the day, and probably, a clock-induced increase in energy metabolism and glial activity is present [55].

Mammalian and insect glial cells modulate circadian neuronal circuitry and behavior via glial calcium signaling [19]. Genetic manipulations of glial vesicle trafficking, the membrane ionic gradient, or internal calcium storage all lead to arrhythmic locomotor activity in Drosophila, an organism in which astrocytes, but not other glial cell types, are relevant for the circadian modulation of behavior. It should be noted that Drosophila and mammalian astrocytes elicit similar functions due to their preserved morphology and molecular signatures. Besides, PER-based glial oscillator is not essential for the free-running behavioral rhythmicity, although the possibility that this oscillator is required for circadian photic sensitivity or the expression of a different rhythm cannot be ruled out [19].

Recently, Xu and colleagues reported the existence of canonical circadian clock genes in mammalian retinal Müller glia. This study not only demonstrated that retinal Müller cells generate molecular circadian rhythms isolated from other retinal cell types but also demonstrated that these retinal cells exhibit unique features of their molecular circadian clock compared to the retina as a complex system. However,
it is important to highlight that the authors mention that both mouse and human Müller cells exhibit species-specific differences in the gene dependence of their clocks [56]. Accordingly, it was observed that human Müller cells exhibit in vitro circadian rhythms in clock gene expression, although the rhythm in these cells does not seem to depend on Per1 expression. Whereas, in mouse Müller cells, knockout or knockdown of Per1 led to arrhythmicity, suggesting that human Müller cells may have a decreased dependence in Per1 expression to regulate rhythmicity [56]. Additional evidence reported by Tosini and Menaker demonstrates that the mammalian neural retina contains a genetically programmed circadian oscillator [57]. Nevertheless, Xu and coworkers propose to Müller glia as a candidate clock cell population in the mammalian retina [56]. The results obtained in both reports indicate that both neurons and glia play an important role in the generation of circadian rhythms in this autonomous oscillator.

2.2. Microglia. These glial cells are the main innate immune cells of the CNS and play essential functions in the maintenance of neuronal circuitry, regulation of behavior, and functional state of neurotransmission [20, 21].

Knowledge about a molecular clock in this type of glial cells is relatively recent. In 2011, it was demonstrated that the clock genes are constitutively expressed in both cultured murine microglia and the microglial cell line BV-2 cells. In the same study, it was also reported that ATP selectively promotes the expression of mRNA and corresponding protein for Per1 via P2X7 purinergic receptor subtype in microglial cells [58]. Years later, it was confirmed that cortical microglia contain an intrinsic molecular clock capable of regulating diurnal changes of its morphological aspect [20]. Specifically, it has been demonstrated in mice that microglia controls the sleep-wake cycle-dependent changes in synaptic strength through the extension and retraction of their processes [21]. Hayashi and colleagues showed that CatS (Cathepsin S, a microglia-specific lysosomal cysteine protease in the brain) exhibits a circadian expression in cortical microglia. Such expression of CatS induces diurnal variations in the synaptic strength of the cortical neurons via the proteolytic modification of the perineuronal environment. Conversely, alterations in CatS lead to hyperlocomotor activity, as well as the deletion of the diurnal variations in the synaptic activity and dendritic spine density of the cortical neurons as a consequence of failure to downscale the synaptic strength during sleep [20, 59]. This process is necessary for the acquisition of subsequent novel information after waking [20]; therefore, dysfunction of microglia intrinsic circadian clock could be involved in social behavior abnormalities [59] and neuropsychiatric disorders, including depression and cognitive impairment [60, 61].

In 2015, Fonken and coworkers reported that microglia possesses circadian clock mechanisms and displays rhythmic fluctuations in both basal inflammatory gene expression and inflammatory potential. It is interesting to note that inflammatory potential in microglia is associated with time-of-day differences, this is because of the circadian differences observed in sickness response [23].

Recently, Nakazato and colleagues demonstrated that Bmal1 modulates Il-6 upregulation in microglial cells exposed to LPS using siRNA targeting Bmal1 and Bmal1-deficient mice [62]. These results suggest that an intrinsic microglial clock may regulate microglial inflammatory responses under pathological conditions in vivo. It was also observed that Bmal1 bindings to the Il-6 promoter region only in cells exposed to LPS; for which, they suggest that histone modification occurred at the Il-6 promoter region with E-box elements [62].

2.3. Oligodendrocytes. These cells are the myelinating glia of the CNS, provide axonal metabolic support [63], and contribute to neuroplasticity [64]. Scarce information regarding clock genes in these cells is available. A previous study suggested that oligodendrocytes’ proliferation depends on the time-of-the-day in the hippocampal hilus, indicating a close connection between the temporal information and glial cells in this structure [65]. To date, there is no report showing that oligodendrocytes have an internal circadian clock. However, it has been suggested that clock genes might regulate OPC (oligodendrocyte precursor cell) proliferation, since these cells in the hippocampus express cyclin D1 [18], which is regulated by Per2 gene [66].

3. Clinical Implications
Recent studies indicate that defective clock genes in glial cells participate in diverse brain pathologies, mainly in psychiatric diseases. However, it is important to keep in mind that a single clock gene can have different repercussions on health and that several clock genes may be related to the same pathology (for detailed review, see reference [67]). Particularly, mutations in Clock, Npas2, and/or Per2 are all involved in a hyperglutamatergic scenario due to a decrease in GLAST expression and as consequence, a reduction in Glu uptake [31, 32, 68]. In this scenario, astrocytic Glu release has clear pathophysiological implications like stroke, multiple sclerosis, and dementia [69]. Additionally, it has been established that Glu regulates the levels of dopamine and other neurotransmitters and neuropeptides that mediate both positive and negative aspects of drug reinforcement and reward. In this manner, both hyper- and hypoglutamatergic states in specific brain areas are directly involved in different stages of addiction, including development, persistence, and abstinence [68]. Fascinatingly, clock genes participate in the modulation of common mechanisms of drug abuse-related behaviors [31, 70].

Moreover, alterations in Per1, Per3, and Bmal1 lead essentially to changes in both short- and long-term memory, chronic oxidative stress in the brain, variations in cocaine sensitization, and association with a number of psychiatric diseases [71–77]. Similarly, Npas2, Gsk3β, Dbp, CRY1, and Clock are involved in variations in drugs sensitization, as well as in diverse psychiatric diseases, mainly bipolar disorder, schizophrenia, Alzheimer, and unipolar major depressive disorder [32, 78–82].

Nowadays, disturbances in the sleep parameters are common. These disturbances are associated with a spectrum of
neurological and psychiatric disorders. Interestingly, clock genes are also involved in variations related with sleep time, sleep fragmentation, and atypical responses following sleep deprivation [83–85]. However, sleep disruptions also have severe consequences in the immune system, leading to an impaired immune function [86, 87]. In line with these reports, it has been established that immune cells exhibit circadian expression of clock genes, which in turn, participate in the regulation of diverse immunological activities. Particularly, REV-ERB is involved in neurodegenerative disorders with an inflammatory component [47]. It has been demonstrated that this clock gene represses macrophage gene expression [88] and targets inflammatory function of macrophages through the direct regulation of Ccl-2 [50]. On the other hand, Bmal1 controls rhythmic trafficking of inflammatory monocytes to sites of inflammation [89].

Table 1: Circadian functions regulated by the glial cells.

<table>
<thead>
<tr>
<th>CG/CCG/molecule</th>
<th>Circadian functions</th>
<th>References</th>
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<tbody>
<tr>
<td><strong>Astrocytes</strong></td>
<td></td>
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<tr>
<td>Clock</td>
<td>Regulation of the glutamatergic system (Glast mRNA and protein levels)</td>
<td>[32]</td>
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<tr>
<td></td>
<td>Modulates ATP release</td>
<td>[55]</td>
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<tr>
<td>Npas2</td>
<td>Regulation of the glutamatergic system (Glast mRNA and protein levels)</td>
<td>[32]</td>
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<td></td>
<td>Regulation of nociceptive processes</td>
<td>[51]</td>
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<tr>
<td>Per1</td>
<td>Modulates ATP release</td>
<td>[55]</td>
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<td></td>
<td>Regulation of the glutamatergic system (GLAST protein levels)</td>
<td>[31, 32]</td>
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<td></td>
<td>Regulation of nociceptive processes</td>
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<tr>
<td>Per2</td>
<td>Modulates ATP release</td>
<td>[55]</td>
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<td></td>
<td>Regulates to cyclin D1</td>
<td>[66]</td>
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<tr>
<td>Bmal1</td>
<td>Modulates the period of the SCN and behavior</td>
<td>[36]</td>
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<td></td>
<td>Regulation of nociceptive processes</td>
<td>[51]</td>
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<tr>
<td>Cry1</td>
<td>Regulation of nociceptive processes</td>
<td>[51]</td>
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<tr>
<td>Gfap</td>
<td>Participates in metabolic exchanges and plasticity</td>
<td>[38, 40–42]</td>
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<tr>
<td>NF-κB</td>
<td>SCN astrocytes mediate the immune signals to the circadian system via NF-κB signaling</td>
<td>[43]</td>
</tr>
<tr>
<td>RORα</td>
<td>Participates in the regulation of the inflammatory response (inhibits NF-κB pathway and regulates IL-6 expression)</td>
<td>[44–46]</td>
</tr>
<tr>
<td>REV-ERBe/REV-ERBβ</td>
<td>Participates in the regulation of the inflammatory response (both isoforms inhibit TNF-induced upregulation of Ccl-2 and Mmp-9; and REV-ERBe inhibits IL-6 upregulation)</td>
<td>[47–50]</td>
</tr>
<tr>
<td>GS</td>
<td>Regulation of the glutamatergic system (glutamate-glutamine metabolic cycle)</td>
<td>[51]</td>
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<tr>
<td>COX-1</td>
<td>Regulation of various spinal sensory functions</td>
<td>[51]</td>
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<tr>
<td>IP3</td>
<td>Modulates ATP release (IP3-dependent calcium signaling)</td>
<td>[55]</td>
</tr>
<tr>
<td>ATP</td>
<td>Regulation of the energy metabolism and glial activity</td>
<td>[55]</td>
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<tr>
<td>Ca^{2+}</td>
<td>Modulation of circadian behavior</td>
<td>[19]</td>
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<td></td>
<td>Regulates the release of gliotransmitters</td>
<td>[35]</td>
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<td></td>
<td>Regulates Bmal1 expression (Glu time- and dose-dependent)</td>
<td>[22]</td>
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<tr>
<td>Glu</td>
<td>Provides the inhibitory astrocytic-neuronal coupling signal during nighttime in the SCN via NMDAR2C</td>
<td>[35]</td>
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<td><strong>Microglia</strong></td>
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<tr>
<td>ATP</td>
<td>Upregulates the Per1 mRNA expression via P2X7 purinergic receptor subtype</td>
<td>[58]</td>
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<tr>
<td>CatS</td>
<td>Regulates the synaptic strength, including neuronal transmission and spine density via the proteolytic modification of the perineuronal environment</td>
<td>[20, 21, 59]</td>
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<tr>
<td>Bmal1</td>
<td>Implicated in the inflammatory response (modulates Il-6 upregulation)</td>
<td>[62]</td>
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<td><strong>Oligodendrocytes</strong></td>
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<tr>
<td>Cyclin D1</td>
<td>Regulation of the OPC proliferation</td>
<td>[18]</td>
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</table>

ATP: adenosine triphosphate; Bmal1: brain and muscle ARNT-like protein 1; Ca^{2+}: calcium; CatS: cathepsin S; CGs: clock genes; CCGs: clock-controlled genes; Ccl-2: monocyte chemoattractant protein-1; Clock: circadian locomotor output cycles kaput; COX-1: cyclooxygenase-1; Cry1: cryptochrome 1; Gfap: glial fibrillary acidic protein; GLAST: glutamate aspartate transporter; Glu: glutamate; GS: glutamine synthetase; Il-6: interleukin-6; IP3: inositol triphosphate; Mmp-9: matrix metalloproteinase-9; NF-κB: nuclear factor-kappaB; NMDAR2C: N-methyl-D-aspartate receptor 2C subunit; Npas2: neuronal PAS domain protein 2; P2X7: purinoreceptor; OPCs: oligodendrocyte precursor cells; Per: period; REV-ERB: reverse Erb; RORα: retinoid-related orphan receptor-alpha; SCN: suprachiasmatic nucleus; TNF: tumor necrosis factor.
these reports together, it is possible to suggest that circadian disruptions exacerbate inflammatory responses in both periphery [90] and CNS [91].

Additionally, it has been shown that RORα is an important molecular player in diverse pathological processes including oxidative stress-induced apoptosis and cerebral hypoxia, both in neurons and astrocytes, due to its neuroprotective properties [44].

Finally, abnormal microglial cells are also associated with neurological disorders [92–94]. Taking into consideration that a risk factor for psychiatric diseases is the dysfunction of the clock system, it is relevant to suggest that the microglial clock might be an interesting target for the development of novel neurological therapeutic agents.

4. Conclusion

The expression of clock genes in glial cells has great importance for the maintenance of a healthy brain (Table 1). Actually, clock genes are relevant for the development of novel strategies for the treatment of a wide range of human diseases such as metabolic and cardiovascular diseases, immune system dysfunction, neuropsychiatric disorders, and even cancer. Specifically, changes in the expression of clock genes in glial cells lead to problems related to an imbalance of the glutamatergic system, resulting in neurological disorders; therefore, understanding the role that glial cells play in brain circadian physiology is extremely relevant.

Conflicts of Interest

The authors declare no competing financial interests.

Acknowledgments

Donaji Chi-Castañeda is supported by SNI-CONACYT, and the work in the lab (Arturo Ortega) is supported by CONACYT-México (255087) and “Soluciones para un México Verde S.A. de C.V.”

References


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