The role of ATP in sleep regulation

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INTRODUCTION: ATP AS A NEUROMODULATOR

Sleep is thought to be essential to replenish energy sources in the brain that are depleted during wakefulness. ATP has long been recognized as an intracellular energy source in the brain and multiple other tissues, and the complex regulation of mitochondrial ATP synthesis is triggered within the human brain in response to daily activities (Kagawa, 2010). ATP has also been proposed as a putative transmitter in sensory nerves, motor nerves, and some CNS neurons (Burnstock, 2007, 2009). In the CNS, ATP signaling can regulate both excitatory and inhibitory neurotransmission. Many neurons can release ATP in an activity-dependent manner, and this molecule can act as a potent neuromodulator for neuron–neuron and neuron–glial signaling (Fields and Stevens, 2000; Fields and Burnstock, 2006; Krueger et al., 2008). ATP and its metabolites regulate multiple processes in the nervous system including sleep regulation (Burnstock et al., 2011).

Recently, it was observed in rats that ATP levels in several brain regions (frontal cortex, basal forebrain, cingulate cortex, and hippocampus) are stable during waking but exhibit a surge during the initial hours of sleep (Dworak et al., 2010). This ATP surge displays a significant positive correlation with the intensity of EEG slow wave activity (SWA; 0.5–4 Hz) in non-rapid eye movement (NREM) sleep, a parameter of sleep depth and homeostatic need for sleep. These results suggest that ATP levels drastically change during sleep in several brain regions and are directly related to SWA in NREM sleep (Dworak et al., 2010). Interestingly, sleep deprivation for 3 h induces a significant reduction in AT P concentration in the frontal cortex and lateral hypothalamus (an area known to predominantly contain wake- and REM-active neurons) although sleep deprivation does not affect the VLPO, an area known to predominantly contain sleep-active neurons (Dworak et al., 2010). Thus, elevations in brain ATP consumption caused by prolonged wakefulness may differ depending on the brain region.

The purpose of this review is to provide an overview of the neurobiological evidence for the role of ATP in sleep regulation. In this review, we will first provide the experimental evidence for an association between adenosine (a product from AMP) and sleep regulation. A presentation of the evidence demonstrating the role of gliotransmission in sleep homeostasis as mediated by astrocytes will follow. Finally, we will address the possibility of an involvement of AMP-activated protein kinase (AMPK), a metabolic sensor, in sleep homeostasis.

ADENOSINE AND SLEEP

The hypnogenic properties of adenosine in specific brain areas have been demonstrated in many studies. Infusion of adenosine and adenosine transport inhibitors into the basal forebrain increases sleep in cats and rats (Portas et al., 1997; Methippara et al., 2005). Extracellular adenosine levels in the basal forebrain measured by microdialysis are lower during NREM and REM sleep in cats, while levels are increased during sleep deprivation and subsequently normalized during recovery sleep (Porkka-Heiskanen et al., 2000). Furthermore, SWA in NREM sleep displays a strong positive correlation with increases in adenosine levels in the basal forebrain and frontal cortex during the recovery period after sleep deprivation (Kalinchuk et al., 2011). These findings suggest that adenosine is a sleep-promoting substance, decreasing the activity of cholinergic wake-promoting neurons in the basal forebrain that are presumably responsible for the accumulation of adenosine during prolonged waking (Porkka-Heiskanen and Kalinchuk, 2011).
An increase in adenosine levels reflects an increased breakdown of cellular ATP (Figure 1). Intracellularly, ATP is metabolized to ADP and then subsequently metabolized to AMP. AMP can be converted back to ATP by adenosine kinase or converted to inosine by adenosine deaminase. In terms of energy state modulation, adenosine kinase and adenosine deaminase are important regulators of intracellular and extracellular adenosine levels. Extracellularly, ATP, ADP, and AMP are converted to adenosine by ecto-5′-nucleotidases. Adenosine concentration between the intra- and extra-cellular spaces is equilibrated by nucleoside transporters. When ATP is broken down in the cell, excess adenosine is transported to the extracellular space by these transporters. The adenosinergic regulation of sleep is predominantly mediated either via A1 receptors (A1Rs) or A2a receptors (A2aRs). The effects of adenosine in the basal forebrain on recovery sleep mentioned above are mostly mediated by A1Rs. Perfusion of an A1R antagonist into the basal forebrain reduces recovery sleep after sleep deprivation, while an A2aR antagonist had no effect on recovery sleep in rats (Gass et al., 2009). However, evidence for a role of A2aRs in the regulation of sleep has also emerged, especially concerning the actions of A2aRs in the leptomeninges under the basal forebrain that are mediated by prostanandin D2 and DP1 receptors (Urade and Hayaishi, 2011). In addition, it was reported that administration of caffeine, a non-selective adenosine receptor antagonist, induces arousal effects in wild-type and A1R-knockout mice, although its effect was not observed in A2aR-knockout mice, indicating that the arousal effects of caffeine are dependent on A2aRs (Huang et al., 2005). Thus, adenosine regulates either the amount of sleep or sleep homeostasis in multiple ways depending on the brain area and receptor subtype.

**GLIA AND SLEEP**

Glia release multiple neuroactive substances that contribute to sleep regulation including ATP, cytokines, and glutamate (Fields and Burnstock, 2006; Krueger et al., 2008). Glia were thought to play a passive and auxiliary role in the brain. However, in the past decade, it has been reported that glia play more active roles in the brain via neuron–astrocyte interactions. Astrocytes can be activated by neurotransmitters released from active neurons they surround. In turn, astrocytes release chemical transmitters to act on neurons pre- and post-synaptically. This process of gliotransmission modulates synaptic activity, with the primary chemicals released from astrocytes being ATP, glutamate, and d-serine, all of which can act to facilitate neural transmission (Hamilton and Attwell, 2010).

Some recent studies have presented evidence that sleep homeostasis depends on gliotransmission and the accumulation of adenosine degraded from ATP. These studies use transgenic mice that express a dominant-negative (dn) SNARE domain selectively in astrocytes, and such mice have impaired gliotransmission (Halassa and Haydon, 2010; Florian et al., 2011). In the dnSNARE mice, SWA in NREM sleep at baseline and during recovery periods after sleep deprivation is attenuated compared with that observed in wild-type mice (Halassa et al., 2009). During sleep deprivation, glutamate released from presynaptic neurons binds...
to postsynaptic AMPA receptors to facilitate excitatory neurotransmission (Figure 2). Glutamate also binds to metabotropic glutamate receptors on the membranes of adjacent astrocytes and can evoke rises in intracellular Ca\textsuperscript{2+} concentration in astrocytes. In turn, astrocytes release ATP when intracellular Ca\textsuperscript{2+} is elevated. ATP released from activated astrocytes is rapidly hydrolyzed to adenosine, which reduces the release of glutamate from the presynaptic terminal by acting on A1 receptors, ultimately resulting in hyperpolarization of the postsynaptic neuron membrane potential (Jones, 2009). The dnSNARE mice, which have an impairment of gliotransmission, display an impaired regulation of sleep homeostasis, suggesting that gliotransmission is essential for sleep pressure accumulation.

It is also well known that sleep loss can induce a deficit in memory task performance in humans and rodents. The dnSNARE mice were also protected against the effects of sleep deprivation in both non-hippocampal- and hippocampal-dependent memory tasks (Halassa et al., 2009; Florian et al., 2011). In addition, i.c.v. injection of the A1R antagonist 8-cyclopentyl-1,3-dimethylxanthine (CPT) into wild-type mice mimicked the phenotype of dnSNARE mice, suggesting that sleep homeostasis and cognitive deficits are due to adenosine actions via A1Rs. Thus, astrocytes regulate sleep pressure, recovery, and associated cognitive consequences mediated by adenosine in terms of dampening neuronal excitability. Interestingly, extracellular ATP also activates purine P2 receptors (e.g. P2X7) on glial cells to release cytokines such as interleukin-1β and tumour-necrosis factor-α, which act as sleep regulatory substances (Krueger et al., 2008, 2010). Mice lacking functional P2X7 receptors have attenuated NREM sleep and SWA in NREM sleep during recovery periods after sleep deprivation compared with wild-type mice (Krueger et al., 2010). These data suggest that extracellular ATP and P2 receptors have an important role in the regulation of sleep homeostasis and link neuro- and gliotransmission to sleep.

**AMPK AND SLEEP**

AMP-activated protein kinase is an efficient sensor of the cellular energy state, and regulates energy balance in the hypothalamus, liver, muscle, adipose tissue, and pancreas (Kahn et al., 2005; Minokoshi et al., 2008). AMPK is activated by phosphorylation when the intracellular AMP/ATP ratio increases, an indication that cellular energy consumed exceeds energy produced. In order to restore energy balance, activated AMPK upregulates catabolic pathways generating ATP such as fatty acid oxidation and glucose uptake, while it downregulates anabolic pathways consuming ATP such as lipid and carbohydrate synthesis. Therefore, dysfunction of peripheral AMPK leads to metabolic syndromes including type 2 diabetes and obesity (Hardie, 2008; Zhang et al., 2009).

Pharmacological experiments have indicated that energy depletion using 2,4-dinitrophenol (DNP), which blocks ATP synthesis and increases the AMP/ATP ratio, increases the amount of NREM sleep and also enhances SWA in NREM sleep (Kalinchuk et al., 2003). In agreement with this study, we have recently demonstrated that i.c.v. injection of 5-aminoimidazole-4-carboxamide riboside (AICAR), an AMP-mimicking activator of AMPK, enhances SWA in NREM sleep, while i.c.v. injection of compound C (CC), an AMPK inhibitor, suppresses SWA compared to vehicle injection in mice (Chikahisa et al., 2009). Furthermore, we have demonstrated that 6 h of sleep deprivation increases AMPK activity in the hypothalamus of mice (Chikahisa et al., 2009). This sleep deprivation-induced activation of AMPK is also observed in the basal forebrain, and is accompanied by decreased ATP levels in
the same brain regions (Dworak et al., 2010). We also observed that mRNA levels of Ca2+/calmodulin (CaM)-dependent protein kinase kinase β (CaMKK2), an additional upstream kinase of AMPK, were increased immediately after 6 h of sleep deprivation and that carnitine palmitoyltransferase 1 (CPT1), a signaling factor downstream of AMPK, was increased during the recovery period after sleep deprivation (Chikahisa et al., 2009). These findings suggest that prolonged wakefulness enhances the activity of central AMPK by increases in the AMP/ATP ratio and/or CaMKK2 expression. Subsequent activation of central AMPK would then cause an enhancement of SWA in NREM sleep (Figure 3).

AMP-activated protein kinase is also known to be involved in circadian rhythms, and directly regulates phosphorylation and degradation of cryptochrome 1 (CRY1), a circadian clock component (Lamia et al., 2009). Furthermore, mice with a genetic disruption of the AMPK pathway show an attenuation of circadian rhythms in the expression of several clock genes in the liver (Lamia et al., 2009). Taken together, AMPK plays an important role in linking energy metabolism and the regulation of circadian and sleep homeostasis. Of course, ATP is itself affected by circadian rhythms in the expression of several clock genes in the liver (Marpegan et al., 2011). Does ATP or its metabolites play a potent physiological role in sleep regulation in the cortex? Does ATP have a role in the connection between hypothalamic and cortical signaling? Further critical research could help understand the pathophysiological mechanisms of sleep disorders and the interaction between sleep impairment and metabolic syndrome is anticipated.

**FUTURE DIRECTIONS**

Due to several advanced studies, an involvement of ATP as an energy source or as a neuromodulator in the regulation of sleep has been identified in the past few years. We now have a better understanding of the functional properties of ATP as a result of individual investigations into the properties of astrocytic, adenosine, and AMPK signaling. However, several questions still remain unanswered. SWA in NREM sleep was recently discovered to be controlled locally in the cortex (Vyazovskiy et al., 2011). Does ATP or its metabolites play a potent physiological role in sleep regulation in the cortex? Does ATP have a role in the connection between hypothalamic and cortical signaling? Further critical research could help understand the pathophysiological mechanisms of sleep disorders and the interaction between sleep impairment and metabolic syndrome is anticipated.

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