Loss of the conserved PKA sites of SIK1 and SIK2 increases sleep need

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Although sleep is one of the most conserved behaviors, the intracellular mechanism regulating sleep/wakefulness remains unknown. We recently identified a protein kinase, SIK3, as a sleep-regulating molecule. Mice that lack a well-conserved protein kinase A (PKA) phosphorylation site, S551, showed longer non-rapid eye movement (NREM) sleep and increased NREMS delta density. S551 of SIK3 is conserved in other members of the SIK family, such as SIK1 (S577) and SIK2 (S587). Here, we examined whether the PKA phosphorylation sites of SIK1 and SIK2 are involved in sleep regulation by generating SIK1S577A and SIK2S587A mice. The homozygous SIK1S577A mice showed a shorter wake time, longer NREMS time, and higher NREMS delta density than the wild-type mice. The heterozygous and homozygous SIK2S587A mice showed increased NREMS delta density. Both the SIK1S577A and SIK2S587A mice exhibited proper homeostatic regulation of sleep need after sleep deprivation. Despite abundant expression of SIK1 in the suprachiasmatic nucleus, the SIK2S587A mice showed normal circadian behavior. Although SIK2 is highly expressed in brown adipose tissue, the male and female SIK2S587A mice that were fed either a chow or high-fat diet showed similar weight gain as the wild-type littermates. These results suggest that PKA-SIK signaling is involved in the regulation of sleep need.

Sleep is a conserved behavior in both vertebrates and invertebrates. However, the molecular mechanism regulating sleep/wakefulness remains to be elucidated. Recently, we identified a kinase, salt-inducible kinase 3 (SIK3), as an important sleep regulator through electroencephalogram (EEG)/electromyogram (EMG)-based sleep screening of randomly mutagenized mice. Mice that express mutant SIK3 lacking the 52 amino acids encoded by exon 13 showed a decrease in wake time and an increase in non-rapid eye movement (NREM) sleep time. Among the 52 amino acids, serine 551 (S551) is a well-conserved protein kinase A (PKA) phosphorylation site and is crucial for the determination of sleep need, suggesting that the PKA-SIK3 pathway is involved in the regulation of sleep/wakefulness.

The SIK family belongs to the AMP-activated protein kinase (AMPK) family of Ser/Thr kinases and is composed of SIK1, SIK2, and SIK3. Similar to other members of the AMPK family, SIKs are primarily activated by LKB1 via phosphorylation of the kinase domain. SIK1 was originally identified as an mRNA induced by a high-salt diet and was named salt-inducible kinase, followed by the in silico identification of SIK2 and SIK3. SIK3 has three well-characterized PKA phosphorylation sites, T469, S551, and S674. Whereas T469 of SIK3 is equivalent to SIK1 T475 and SIK2 T484, S551 of SIK3 is equivalent to SIK1 S577 and SIK2 S587. Phosphorylation of these PKA-phosphorylation sites promotes 14-3-3 binding to SIKs, which leads to changes in subcellular localization and kinase activity. Since SIK3S551A mice showed longer NREM sleep and higher NREMS delta density than wild-type mice, we focused on S551-equivalent PKA-phosphorylation sites, S577 of SIK1 and S587 of SIK2, hypothesizing that these PKA-phosphorylation sites are involved in sleep/wake regulation similar to S551 of SIK3.

SIK1 and SIK2 have been reported to regulate circadian behavior and energy metabolism. The knockdown of SIK1 in the suprachiasmatic nucleus (SCN) resulted in a rapid phase shift of the circadian rhythm, in other words, resistance to jet lag. Mice deficient in Sik1 showed lower body weights under both chow and high-fat diets than...
Figure 1. SIK family mRNA expression and mutant proteins. (a) Scheme of SIK1, SIK2, and SIK3. The serine residue in the PKA consensus sequence is conserved among the family. Although there are multiple protein isoforms of SIK3, this scheme shows the longest isoform. (b) Digital PCR results of \( \text{Sik1} \), \( \text{Sik2} \) and \( \text{Sik3} \) mRNA of the cerebral cortex, hippocampus, hypothalamus, liver and brown adipose tissue (BAT) of the wild-type mice (\( n = 4 \)). Each sample was measured in duplicate. One-way analysis of variance followed by Tukey’s test. (c-e) \text{In situ} hybridization of \( \text{Sik1} \) and \( \text{Sik2} \). (c) \( \text{Sik1} \) mRNA was strongly expressed in the suprachiasmatic nucleus (SCN) and broadly expressed in the forebrain. Scale bar, 500 μm. (d) \text{In situ} hybridization showed that \( \text{Sik2} \) mRNA was broadly expressed in the forebrain. Scale bar, 500 μm. (e) \( \text{Sik1} \) and \( \text{Sik2} \) were expressed in the hippocampal dentate gyrus of the wild-type mice (upper and middle). \( \text{Sik2} \) expression was not detected in the dentate gyrus of the \( \text{Sik2} \)-deficient mice (lower). Scale bars, 100 μm. (f-h) \( \text{Sik1} \) (f), \( \text{Sik2} \) (g) and \( \text{Sik3} \) (h) mRNA expression in the cerebral cortex, hypothalamus, BAT, liver and adrenal gland after one week of high-salt diet feeding. Two-tailed t-test with Bonferroni correction. (i) \( \text{SIK2} \) protein was expressed in the BAT of the \( \text{Sik2}^{+/+} \) and \( \text{Sik2}^{S587A/S587A} \) mice. \( \text{SIK2} \) was not detected in the BAT of the \( \text{Sik2} \)-deficient mice. (j) FLAG-\( \text{SIK1} \) WT, FLAG-\( \text{SIK1} \) S577A, FLAG-\( \text{SIK2} \) WT, FLAG-\( \text{SIK2} \) S587A, FLAG-\( \text{SIK3} \) WT, and FLAG-\( \text{SIK3} \) S551A transiently expressed in HEK293 cells were immunoprecipitated with an anti-FLAG antibody and then subjected to immunoblotting using.
The SIK1 S577A and SIK2 S587A proteins did not bind to 14-3-3. Since SIK1 was originally identified as a molecule induced by a high-salt diet in the rat adrenal gland, we examined whether a high-salt diet affects SIK1, 2 and 3 expression in the brain, BAT, liver and adrenal gland. One week of a high-salt diet did not affect the SIK1 mRNA expression in the cerebral cortex, hypothalamus, BAT, liver or adrenal gland (Fig. 1f). A high-salt diet increased the SIK2 mRNA expression in the adrenal gland (Fig. 1g) and did not cause significant changes in the SIK3 mRNA expression in all tissues examined (Fig. 1h). We also confirmed the SIK2 protein expression in the BAT (Fig. 1i, S1a,c) and the brain (Fig. S1b,d).

The SIK1 S577A and SIK2 S587A proteins did not bind to 14-3-3. For characterization of the SIK1 S577A and SIK2 S587A proteins, FLAG-tagged SIK protein variants were transiently expressed in HEK293 cells. Since SIKs have a RRAS motif, a consensus sequence for PKA, immunopurified FLAG-tagged SIK1, 2, and 3 were detected with an anti-phospho-PKA substrate motif antibody (Fig. 1i). In contrast, the FLAG-tagged SIK1 S577A, SIK2 S587A, and SIK3 S551A showed only faint immunoreactivity to the anti-phospho-PKA substrate motif antibody. The 14–3–3 protein was purified with the FLAG-tagged SIK1, 2, and 3 but not with the FLAG-tagged SIK1 S577A, SIK2 S587A and SIK3 S551A (Fig. 1j), as previously reported.

The SIK1 S577A mice showed an increased sleep need. We generated SIK1 S577A mice to examine whether the presence of the PKA phosphorylation site, S577, of SIK1 is required for normal sleep/wakefulness (Fig. 1a). The external appearance, such as coat color, whisker morphology and eye shape, of the heterozygous and homozygous SIK1 S577A mice was indistinguishable from that of the SIK1 +/− mice. The heterozygous and homozygous SIK1 S577A mice did not show any abnormal locomotion or sensory response against touch or sound. The heterozygous and homozygous SIK1 S577A mice were fertile. The male and female SIK1 S577A and SIK1 S577A mice showed similar weights compared to the wild-type littermates at the age of 8 weeks (Fig. S1a).

The SIK1−/− mice had a higher delta range power during REMS than the wild-type littermates (Fig. 2k). There was no significant difference in the delta range power during NREMS among the genotypic groups (Fig. 2l). The SIK1−/− mice had a higher delta range power during REMS than the wild-type littermates (Fig. 2m). Thus, the SIK1−/− homozygous mice showed increased sleep need in terms of time and EEG spectrum.
To investigate the homeostatic regulation of sleep/wakefulness, we performed sleep deprivation for 6 h with gentle handling. There was no difference in time spent in each vigilance state during the light phase after sleep deprivation among the genotypic groups (Fig. 3a–c). The Sik1S577A/S577A mice showed a shorter wake time during the dark phase than the Sik1 S577A/+ mice, and the Sik1S577A/+ mice showed a shorter wake time than the Sik1+/+ mice (Fig. 3a). Consistently, the Sik1S577A/S577A mice showed longer NREMS time during the dark phase than the Sik1S577A/+ and Sik1+/+ mice (Fig. 3b). The Sik1S577A/+ mice showed longer REMS time during the dark phase than the Sik1+/+ mice (Fig. 3c). The change in NREMS delta power for 2 h after sleep deprivation was higher than that under the basal condition in all genotypic groups (Fig. 3d). Sleep deprivation increased the NREMS delta power.

Figure 2. Sleep/wake behavior of the Sik1S577A mice. (a–c) Total time spent in wakefulness (a), NREMS (b) and REMS (c) of the Sik1+/+, Sik1S577A/+ and Sik1S577A/S577A mice. (d–f) Episode duration of wakefulness (d), NREMS (e) and REMS (f) of the Sik1+/+, Sik1S577A/+ and Sik1S577A/S577A mice. (g–i) Circadian variation of wakefulness (g), NREMS (h) and REMS (i) of the Sik1+/+, Sik1S577A/+ and Sik1S577A/S577A mice. (j) Hourly NREMS delta power density over 24 h. (k–m) EEG power density of the Sik1+/+, Sik1S577A/+ and Sik1S577A/S577A mice during wakefulness (k), NREMS (l) and REMS (m). 13 male mice for each group. One-way analysis of variance (ANOVA) followed by Tukey's test (a–f, k–m). One-way repeated-measures ANOVA followed by Tukey's test (j). Data are presented as the mean ± SEM.
to a similar extent among the genotypic groups (Fig. 3e). The Sik1S577A mice showed a higher NREMS delta density during the light phase than the Sik1 mice (Fig. 3f). Similar to basal sleep/wakefulness, the Sik1S577A/S577A mice had a higher delta power during wakefulness than the wild-type littermates after sleep deprivation. Figure 3. Sleep/wakefulness of the Sik1S577A mice after sleep deprivation. (a–c) Hourly time spent in wakefulness (a), NREMS (b), and REMS (c) of the Sik1+/+, Sik1S577A/+, and Sik1S577A/S577A mice after 6-h sleep deprivation. (d) NREMS delta power of the Sik1S577A mice under the basal state and for two hours after 6-h sleep deprivation. (e) NREMS delta density change of the Sik1S577A mice during ZT6-7, ZT8-9, and ZT10-11 h after 6-h sleep deprivation. (f) Hourly NREMS delta power density of the Sik1S577A mice after 6-h sleep deprivation. (g–i) EEG power spectra of the Sik1S577A mice during two hours after 6-h sleep deprivation during wakefulness (g), NREMS (h), and REMS (i). One-way repeated-measures analysis of variance (ANOVA) followed by Tukey’s test (a–c,f). Two-tailed paired T-test (d). Two-way ANOVA (e). One-way ANOVA followed by Tukey’s test (g–i). 13 male mice for each group. Data are presented as the mean ± SEM.
deprivation (Fig. 3g), and there was no significant difference in the delta range power during NREMS among the genotypic groups (Fig. 3h). The Sik1S577A/S577A mice had a higher delta range power during REMS than the wild-type littermates (Fig. 3i). These results suggest normal homeostatic sleep regulation in the Sik1S577A mice.

Next, we generated Sik2S587A mice to investigate the role of the phosphorylation of SIK2 S587 in sleep/wake behavior (Fig. 1a). We confirmed that a SIK2 protein of the same size as wild-type protein was expressed in the homozygous Sik2S587A mice (Fig. 1i). The SIK2 levels in the brains and BAT of the Sik2S587A/S587A mice were lower than those of the Sik2+/+ mice (Fig. S1). The external appearance of the heterozygous and homozygous Sik2S587A mice was indistinguishable from that of the Sik2+/+ mice. Although there was no significant difference in body weight at the age of 8 weeks of the male mice with the Sik2S587A genotype (Fig. S2b), the body weight of the female heterozygous and homozygous Sik2S587A mice was lower than that of the wild-type littermates. The heterozygous and homozygous Sik2S587A mice did not show any abnormal locomotion or sensory response. The heterozygous and homozygous Sik2S587A mice were fertile.

Both the Sik2S587A/+ and Sik2S587A/S587A mice had a total time spent in each vigilance state similar to that of the wild-type mice (Fig. 4a–c). There was no significant difference in episode duration of each vigilance state among the genotypic groups (Fig. 4d–f). EEG spectral analysis revealed a higher NREMS delta density of the Sik2S587A/+ and Sik2S587A/S587A mice than that of the Sik2+/+ mice throughout the entire day (Fig. 4g). There was no difference in the delta range power in each vigilance state among the genotypic groups (Fig. 4h–j). Thus, the Sik2S587A homozygous mice showed increased sleep need in terms of EEG spectrum.

After sleep deprivation for 6 h, the amounts of time spent in wakefulness, NREMS and REMS were similar during the light and dark phases (Fig. 5a–c). The change in NREMS delta power over 2 h after sleep deprivation was higher than that under the basal condition in all genotypic groups (Fig. 5d). Sleep deprivation increased the NREMS delta power to a similar extent among the genotypic groups (Fig. 5e). Similar to basal sleep/wakefulness, the Sik2S587A/+ and Sik2S587A/S587A mice showed a higher NREMS delta density during the light and dark phases than the Sik2+/+ mice (Fig. 5f). The Sik2S587A/+ and Sik2S587A/S587A mice showed higher

Figure 4. Sleep/wake behavior of the Sik2S587A mice. (a–c) Total time spent in wakefulness (a), NREMS (b) and REMS (c) of the male Sik2+/+, Sik2S587A/+, and Sik2S587A/S587A mice. (e–f) Episode duration of wakefulness (d), NREMS (e) and REMS (f) of the Sik2+/+, Sik2S587A/+, and Sik2S587A/S587A mice. (g) Hourly NREMS delta power density over 24 h. (h–j) EEG power density of the Sik2+/+, Sik2S587A/+, and Sik2S587A/S587A mice during wakefulness (h), NREMS (i) and REMS (j). Sik2+/+, n = 12; Sik2S587A/+, n = 13; Sik2S587A/S587A, n = 12. One-way analysis of variance (ANOVA) followed by Tukey’s test (a–f, h–j). One-way repeated-measures ANOVA followed by Tukey’s test (g). Data are presented as the mean ± SEM.

**The Sik2S587A mice showed a mild increase in sleep need.** Next, we generated Sik2S587A mice to investigate the role of the phosphorylation of SIK2 S587 in sleep/wake behavior (Fig. 1a). We confirmed that a SIK2 protein of the same size as wild-type protein was expressed in the homozygous Sik2S587A mice (Fig. 1i). The SIK2 levels in the brains and BAT of the Sik2S587A/S587A mice were lower than those of the Sik2+/+ mice (Fig. S1). The external appearance of the heterozygous and homozygous Sik2S587A mice was indistinguishable from that of the Sik2+/+ mice. Although there was no significant difference in body weight at the age of 8 weeks of the male mice with the Sik2S587A genotype (Fig. S2b), the body weight of the female heterozygous and homozygous Sik2S587A mice was lower than that of the wild-type littermates. The heterozygous and homozygous Sik2S587A mice did not show any abnormal locomotion or sensory response. The heterozygous and homozygous Sik2S587A mice were fertile.

Both the Sik2S587A/+ and Sik2S587A/S587A mice had a total time spent in each vigilance state similar to that of the wild-type mice (Fig. 4a–c). There was no significant difference in episode duration of each vigilance state among the genotypic groups (Fig. 4d–f). EEG spectral analysis revealed a higher NREMS delta density of the Sik2S587A/+ and Sik2S587A/S587A mice than that of the Sik2+/+ mice throughout the entire day (Fig. 4g). There was no difference in the delta range power in each vigilance state among the genotypic groups (Fig. 4h–j). Thus, the Sik2S587A homozygous mice showed increased sleep need in terms of EEG spectrum.

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Figure 5. Sleep/wakefulness of the Sik2^{S587A} mice after sleep deprivation. (a–c) Hourly time spent in wakefulness (a), NREMS (b), REMS (c) of the male Sik2^{+/+}, Sik2^{S587A/+}, and Sik2^{S587A/S587A} mice. (d) NREMS delta power of the Sik2^{S587A} mice under the basal state and for 2 h after 6-h sleep deprivation. (e) NREMS delta density change during ZT6-7, ZT8-9, and ZT10-11 h after 6-h sleep deprivation. (f) Hourly NREMS delta power density of the Sik2^{S587A} mice after 6-h sleep deprivation. (g–i) EEG power spectra of the Sik2^{S587A} mice during two hours after 6-h sleep deprivation during wakefulness (g), NREMS (h), REMS (i). Sik2^{+/+}, n = 9; Sik2^{S587A/+}, n = 9; Sik2^{S587A/S587A}, n = 10. One-way repeated-measures analysis of variance (ANOVA) followed by Tukey’s test (a–c, f). Two-tailed paired T-test (d). Two-way ANOVA (e). One-way ANOVA followed by Tukey’s test (g–i). Data are presented as the mean ± SEM.
delta range power during wakefulness than the Sik2\(^{S587A}\) mice (Fig. 5g). There was no difference in the delta range power in NREMS and REMS among the genotypic groups (Fig. 5h,i).

Normal circadian behavior of the Sik1\(^{S577A}\) mice. Since Sik1 is expressed in the SCN (Fig. 1c) and was reported to be involved in the phase shift of circadian rhythm based on deficiency studies, we examined the circadian behavior of the Sik1\(^{S577A}\) mice. Under constant darkness, the Sik1\(^{S577A/+}\) and Sik1\(^{S577A/S577A}\) mice showed a circadian period similar to that of the wild-type littermates (Fig. 6a–c). Furthermore, the Sik1\(^{S577A/+}\) and Sik1\(^{S577A/S577A}\) mice showed re-entrainment to a new circadian time with a similar phase shift to the wild-type mice (Fig. 6d–f). Thus, Sik1\(^{S577A}\) mutant mice showed normal circadian behavior in terms of endogenous circadian period length and re-entrainment.

Normal body weight growth of the Sik2\(^{S587A}\) mice. Since Sik2 is highly expressed in BAT (Fig. 1b), which regulates energy metabolism, we examined the body weight growth of the Sik2\(^{S587A}\) mice. At the age of 6 weeks, the body weights of the male Sik2\(^{S587A}\) mice fed a chow or a high-fat diet were similar to those of the wild-type littermates (Fig. 7a). The body weights of the female Sik2\(^{S587A/S587A}\) mice on chow were smaller than those of the Sik2\(^{+/+}\) mice (Fig. 7b). However, there was no difference between the female Sik2\(^{S587A/S587A}\) mice on a high-fat diet (Fig. 7b). At the age of 20 weeks, both male and female Sik2\(^{S587A}\) mice fed a chow or a high-fat diet showed body weights similar to those of their wild-type littermates (Fig. 7c,d). There was no significant difference in the body weight gain from 6 weeks to 20 weeks of age among the genotypic groups for both sexes (Fig. 7e,f). Thus, male and female Sik2\(^{S587A}\) mice have normal body weight regulation.

Discussion

In this study, we examined the sleep/wake behavior of mice lacking conserved PKA phosphorylation sites of SIK1 and SIK2. Both the Sik1\(^{S577A}\) mice and the Sik2\(^{S587A}\) mice showed an increased NREMS delta density, an index of sleep, which is consistent with the results of the Sik3\(^{S551A}\) mice that lack an exon 13-encoded region and the Sik3\(^{S551A}\) mice that lack a conserved PKA phosphorylation site. The homozygous Sik1\(^{S577A}\) mice showed a shorter total wake time and longer total NREMS time than the wild-type mice in addition to an increased NREMS delta density, an index of sleep, which is consistent with the results of the Sik3\(^{S551A}\) mice that lack an exon 13-encoded region and the Sik3\(^{S551A}\) mice that lack a conserved PKA phosphorylation site. The homozygous Sik1\(^{S577A}\) mice showed a shorter total wake time and longer total NREMS time than the wild-type mice in addition to an increased NREMS delta density, an index of sleep, which is consistent with the results of the Sik3\(^{S551A}\) mice that lack an exon 13-encoded region and the Sik3\(^{S551A}\) mice that lack a conserved PKA phosphorylation site. The homozygous Sik1\(^{S577A}\) mice showed a shorter total wake time and longer total NREMS time than the wild-type mice in addition to an increased NREMS delta density, an index of sleep, which is consistent with the results of the Sik3\(^{S551A}\) mice that lack an exon 13-encoded region and the Sik3\(^{S551A}\) mice that lack a conserved PKA phosphorylation site. The homozygous Sik1\(^{S577A}\) mice showed a shorter total wake time and longer total NREMS time than the wild-type mice in addition to an increased NREMS delta density, an index of sleep, which is consistent with the results of the Sik3\(^{S551A}\) mice that lack an exon 13-encoded region and the Sik3\(^{S551A}\) mice that lack a conserved PKA phosphorylation site. The homozygous Sik1\(^{S577A}\) mice showed a shorter total wake time and longer total NREMS time than the wild-type mice in addition to an increased NREMS delta density, an index of sleep, which is consistent with the results of the Sik3\(^{S551A}\) mice that lack an exon 13-encoded region and the Sik3\(^{S551A}\) mice that lack a conserved PKA phosphorylation site. The homozygous Sik1\(^{S577A}\) mice showed a shorter total wake time and longer total NREMS time than the wild-type mice in addition to an increased NREMS delta density, an index of sleep, which is consistent with the results of the Sik3\(^{S551A}\) mice that lack an exon 13-encoded region and the Sik3\(^{S551A}\) mice that lack a conserved PKA phosphorylation site.
littermates. Sleep deprivation increased the NREMS delta density to a similar extent, suggesting that the Sik1S577A mice showed proper homeostatic sleep regulation. Both the heterozygous and homozygous Sik2 S587A mice exhibited an increased NREMS delta density, but the Sik2S587A mice did not alter the time spent in wakefulness and sleep. Lower brain expression of Sik2 than Sik3 accounts for the milder sleep phenotype of the Sik2 S587A mice than that of the Sik3 S551A mice. Differential expression patterns between Sik2 and Sik3 may cause normal sleep time of the Sik2 S587A mice. Thus, the current results as a whole support the idea that PKA-SIK signaling is involved in the regulation of sleep need. Importantly, KIN-29, the Caenorhabditis elegans homologue of SIK is involved in the regulation of sleep1,25. Consistent with previous studies12,16,11, the Sik1 S577A, Sik2 S587A, and Sik3 S551A proteins showed reduced immunoreactivities for phospho-PKA substrate motifs. The remaining immunoreactivities for the phospho-PKA motif of Sik1 S577A, Sik2 S587A, and Sik3 S551A are attributed to other PKA recognition sites, such as Sik1 T475, Sik2 T484 and Sik3 T46911. In addition, the Sik1 S577A, Sik2 S587A, and Sik3 S551A proteins showed decreased binding to 14-3-3, as previously reported12,16,11. Importantly, each PKA phosphorylation site of SIKs has differential effects on binding to 14-3-3. Berggreen et al. showed much lower 14-3-3 binding of the Sik3 T469A and Sik3 S551A proteins than the wild-type Sik3 using HEK293 cells10. Sonntag et al. showed that the alanine substitution of Sik2 at S358 or S387 led to a similar decrease in 14-3-3 binding in HEK 293 cells11, which

Figure 7. Body weight growth of the Sik2 S587A mice fed chow or a high-fat diet. (a) Body weights of the male Sik2 S587A mice at the age of 6 weeks. (b) Body weights of the female Sik2 S587A mice at the age of 6 weeks. (c) Body weights of the male Sik2 S587A mice at the age of 20 weeks. (d) Body weights of the female Sik2 S587A mice at the age of 20 weeks. (e) Body weight gain of the male Sik2 S587A mice from 6 weeks to 20 weeks of age. (f) Body weight gain of the female Sik2 S587A mice from 6 weeks to 20 weeks of age. Two-way analysis of variance followed by Sidak’s test. All data are presented as the mean ± SEM. Male Sik2 S587A/+ mice (chow n = 15, HFD n = 11). Male Sik2 S587A/+ mice (chow n = 16, HFD n = 9). Male Sik2 S587A/S587A mice (chow n = 6, HFD n = 7). Female Sik2 S587A/+ mice (chow n = 12, HFD n = 12). Female Sik2 S587A/S587A mice (chow n = 16, HFD n = 15). Female Sik2 S587A/S587A mice (chow n = 15, HFD n = 9).
is consistent with the decreased 14-3-3 binding observed in the present study. However, Henriksson et al. showed that the alanine substitution of SS43 or SS58 of SIK2 decreased 14-3-3 binding, but the alanine substitution of SIK2 SS67 did not affect binding to 14-3-3 in HEK293 cells. Although the reason for this discrepancy is unknown, the effect of PKA on SIKs may be complex depending on the context and the presence of other upstream kinases.

The phosphorylation of the PKA site of SIK1 regulates the nucleo-cytoplasmic localization of SIK1, whereas SIK2 and SIK3 remain localized in the cytoplasm. Regardless of subcellular localization, it was reported that SIKs commonly phosphorylate the same substrates. CAMP-responsive element-binding protein (CREB)-regulated transcription coactivators (CRTC) are the best characterized substrates of SIKs, class IIA histone deacetylases (HDACs) have also been reported as SIK substrates. Phosphorylation of SIKs by PKA prevents SIKs from phosphorylating substrates through 14-3-3 binding and/or nucleocytoplasmic translocation, which subsequently alters gene expression that may lead to altered phosphorylation of sleep need-index phosphoproteins (SNIPPs).

Consistent with a previous report indicating that SIK1 in the SCN is necessary for proper circadian re-entrainment, SIK1 is highly expressed in the SCN. However, the SIK2-/- mice showed normal circadian rhythm and re-entrainment to an advanced light-dark cycle, suggesting that the functional change of SIK1 has little effect on circadian behavior. Similarly, the SIK3-/- mice showed an enhanced sleep need and normal circadian behavior, while SIK3 deficiency caused abnormal circadian behavior in mice and fruit flies. However, since systemic SIK3 deficiency caused severe growth retardation and metabolic and skeletal abnormalities, the neuron-specific SIK3 deficiency is necessary to examine the role of brain SIK3 in circadian behavior in detail.

Male SIK2+/-, 23,31. SIK2-/- mice exhibited body weights similar to those of the wild-type littermates fed either a high-fat diet or chow. Although the female SIK2+/-, mice showed lower body weights than the wild-type controls at the ages of 6 and 8 weeks, they showed a normal body weight gain as they grew. These results are not consistent with a previous report demonstrating that mice with specific SIK2 S587A overexpression in BAT showed high-fat diet-induced obesity. Thus, the SIK2 S587A protein in the BAT may promote weight gain, but SIK2 S587A outside the BAT may counteract the effect on the BAT. Although we did not examine this issue in our study, the SIK2 S587A mice may show altered glucose metabolism since higher blood glucose was found in SIK2-deficient mice and those with pancreatic beta cell specific SIK2 ablation, but not liver-specific SIK2 deficiency, than wild-type mice. Although the mice deficient in SIK1 showed lower body weights under both chow and high-fat diets, the male and female SIK1-/- mice did not show any significant change in body weight. Thus, PKA-SIK1/2 signaling does not play a major role in energy metabolism.

We also examined whether a high-salt diet affects SIK expression. Although a high-salt diet for seven days strongly induced SIK1 mRNA in the adrenal gland of rats, it did not observe a significant increase in SIK1 mRNA in the adrenal gland of mice. This unexpected result may be due to species differences and/or some technical differences. Instead, we found that a high-salt diet increased the SIK2 mRNA levels in the adrenal gland and liver. SIK1 mRNA showed a higher individual difference than SIK2 and SIK3 mRNA.

One limitation of our study is the protein level characterization of mutant mice. Although several antibodies for SIK1 are commercially available, all the SIK1 antibodies we tested did not work. Therefore, we generated polyclonal antibodies in rabbits for SIK1. However, our SIK1 antibodies are not sensitive enough to detect endogenous SIK1, which may be expressed at a low level according to the digital PCR results. For SIK2, a good anti-SIK2 antibody (Cell Signaling Technology #6919) detected SIK2 in the BAT of the wild-type mice, and the corresponding bands were not detected in the SIK2-deficient mice (Fig. S4a). However, this SIK2 antibody showed a band similar in size to SIK2 in brains, which was found for both the wild-type and SIK2 KO mice (Fig. S4b). Then, we used a different antibody (Merck #07-1378), which detects SIK2 in brains (Fig. S4d) and BAT (Fig. S4c). Although identification of endogenous proteins interacting with SIKs in the brain is crucial for the mechanistic understanding of how the SIK family regulates sleep need, the low abundance of SIK2 protein in the brain makes immunoprecipitation difficult. This issue needs to be addressed in future studies using mice that have endogenous SIKs with epitope tags.

We found that SIK2 (SS587A) protein level in the brains of the SIK2 S857/857A mice was lower than the level of SIK2 protein in the wild-type mice. Additionally, the amount of SIK2 (SS587A) protein in BAT of the SIK2 S587/857A mice was lower than the amount of SIK2 protein in the wild-type mice. This result suggests that phosphorylation at SS587A affects the protein amount and/or stability. Since 14-3-3 has been reported to stabilize the binding protein, reduced binding of SIK2 SS587A to 14-3-3 may result in reduced SIK2 SS587A levels. Thus, PKA may regulate SIK2 in multiple aspects, including kinase activity and protein stability, via 14-3-3 binding, which leads to altered phosphorylation of substrates and sleep need.

In summary, the current study broadened and confirmed the hypothesis that PKA-SIK signaling constitutes an intracellular signaling pathway regulating sleep/wakefulness, as was originally conceived based on findings in SIK3 mutant mice. Elucidation of downstream signals is a challenge for the near future.

Methods

Animals. All animal experiments were conducted in accordance with the Guidelines for Animal Experiments of the University of Tsukuba and were approved by the Institutional Animal Care and Use Committee of the University of Tsukuba (approved protocol ID #1800094). SIK2-/- and SIK2 S587A mice and littermate wild-type mice were maintained on the C57BL/6 background. Male C57BL/6 mice were obtained from CLEA Japan. SIK2-deficient mice (#nbio071) were obtained from the Animal Resource Bank of National Institutes of Biomedical Innovation, Health and Nutrition, Japan. Mice were housed under humidity- and temperature-controlled conditions on a 12:12 h light/dark cycle. Food was provided ad libitum. Mice were weaned at 4 weeks of age and were housed in groups of four or five.
**Generation of the Sik1<sup>SS7A</sup> and Sik2<sup>SS8A</sup> mice.** We created Sik1<sup>SS7A</sup> and Sik2<sup>SS8A</sup> mice as previously reported with some modifications<sup>40</sup>. In brief, to produce the Sik1<sup>SS7A</sup> and Sik2<sup>SS8A</sup> mice, we used Cas9 protein, guide RNA (gRNA) and donor oligonucleotide. The gRNAs synthesized using the GeneArt Precision gRNA Synthesis Kit (ThermoFisher), were 5′-TCTGATACCCCTCTACTCA-3′ for Sik1<sup>SS7A</sup> and 5′-TCAGATGCTCCCTTACCA-3′ for Sik2<sup>SS8A</sup>. The donor oligonucleotides for Sik1 and Sik2 were 151- and 113-bases (Integrated DNA Technologies), respectively, which have a sequence to introduce a serine-to-alanine substitution and several silent mutations to disrupt the PAM sequence. Pregnant mare serum gonadotropin and human chorionic gonadotropin were intraperitoneally injected into female C57BL/6 N mice (Charles River Laboratories), and unfertilized oocytes were collected from their oviducts. We then performed in vitro fertilization with these oocytes and sperm from male C57BL/6 N mice. The gRNA, donor oligonucleotide, and GeneArt Platinum Cas9 Nuclease (ThermoFisher) were electroporated into zygotes using a NEPA 21 electroporator (NEPAGNE). After electroporation, embryos developed to the morula stage in KSOM-AA medium. The next day, the electroporated embryos, which developed to the two-cell stage, were transferred into pseudopregnant ICR mice. The genomic DNA sequence of F0 mice was examined by direct sequencing. For routine genotyping, we used the dCAPS method<sup>41</sup> using primers designed with dCAPS Finder 2.0 (http://helix.wustl.edu/dcaps/dcaps.html). For Sik1<sup>SS7A</sup> genotyping, the genomic DNA was amplified using Sik1-Fw (5′-TCTGGA GCAGACGAGTTA-3′) and Sik1-Rv (5′-CTGACAGGCTGGGATA-3′) to produce a 190-bp fragment and then digested with Ddel to produce 70-bp and 120-bp fragments when the allele was wild-type. For Sik2<sup>SS8A</sup> genotyping, the genomic DNA was amplified using Sik2-Fw (5′-TCCCCCGTGACGTCGAGAAAGCCGACAGAA-3′) and Sik2-Rv (5′-CTTTGGTCTGAGATTGTCG-3′) to produce a 215-bp fragment and then digested with HinfI to produce 180-bp and 30-bp fragments when the allele was wild-type. F0 mice were mated with wild-type C57BL/6 mice to obtain heterozygous F1 offspring. Behavioral analysis was performed using F1 or later generations.

**High-salt diet feeding.** For a high-salt diet, NaCl was added to MF (containing 0.3% NaCl, Oriental Yeast) to a concentration of 6.6% NaCl by weight<sup>13-15</sup>, and then water was added to mix NaCl and MF. Mice were fed a high-salt diet or an MF Chow diet for one week. One week of feeding with a high-salt diet did not affect the body weight but increased water intake and urination.

**EEG/EMG electrode implantation surgery.** At 8–12 weeks of age, male mice were implanted with EEG/EMG electrodes with 4 EEG electrode pins and 2 flexible stainless EMG wires under anesthesia using isoflu- rane (4% for induction, 2.5% for maintenance)<sup>46</sup>. The EEG electrode pins were positioned over the frontal and occipital cortices (anteroposterior (AP): 0.5 mm, mediolateral (ML): 1.3 mm, dorsoventral (DV): – 1.3 mm and AP: – 4.5 mm, ML: 1.3 mm, DV: – 1.3 mm) and attached to the skull using dental cement. Then, the EMG wires were inserted into the neck muscles. All mice were allowed at least 4 to 7 days for recovery from the recording surgery. After the recovery period, all mice were attached to a tether cable and then allowed to habituate to the recording conditions for 7 days.

**EEG/EMG recording and analysis.** EEG/EMG recordings were analyzed as previously described<sup>46</sup>. The recording room was kept under a 12:12 h light/dark cycle and a constant temperature (24–25 °C). The EEG/EMG data were visualized and semiautomatically analyzed by MATLAB-based software. The sleep/wake state in each 20-s epoch was classified as NREMS, REMS or wakefulness. Wakefulness was scored based on the criteria of the presence of fast EEG, high amplitude and variable EMG. NREMS was staged based on high amplitude, delta (1–4 Hz) frequency EEG and low EMG tonus. REMS was characterized by theta (6–9 Hz)-dominant EEG and EMG atonia. For analysis of baseline sleep/wake behavior, EEG/EMG signals were recorded for 2 consecutive days from the onset of the light phase, ZT0. The total amounts of time spent in wakefulness, NREMS and REMS were derived by summing the total number of 20-s epochs in each state. Mean episode durations were determined by dividing the total time spent in each state by the number of episodes of that state. EEG signals were subjected to fast Fourier transform analysis from 1 to 30 Hz with 1–Hz bins using MATLAB-based custom software. For normalization of the EEG power density, the EEG power density in each frequency bin was expressed as a percentage of the mean EEG power in a relatively constant higher frequency range (16–30 Hz)<sup>2</sup>. The hourly delta density was calculated as the hourly averages of delta power (1–4 Hz) to total EEG power density (1–4 Hz) to total EEG power (1–30 Hz) at each 20-s epoch or all epochs. For sleep deprivation, the mice were sleep-deprived for 6–8 h during NREMS indicates the hourly averages of delta density, which is the ratio of delta power (1–4 Hz) to total EEG power (1–30 Hz). The hourly delta density during NREMS indicates the hourly averages of delta density, which is the ratio of delta power (1–4 Hz) to total EEG power (1–30 Hz) at each 20-s epoch or all epochs. For sleep deprivation, the mice were sleep-deprived for six hours from the onset of light phase ZT0 by gentle handling<sup>47</sup>. During that time, food and water were available. For evaluation of the effect of sleep deprivation, NREMS delta power during the first hour after sleep deprivation was expressed relative to the same ZT of the basal recording or relative to the mean of the basal recording.

**In situ hybridization.** In situ hybridization was performed as previously described with some modifications<sup>46</sup>. Fragments of the coding regions for Sik1 and Sik2 were PCR amplified using Sik1-Fw (5′-ATAGA CTGTG ATCTC CACAG CTCAC TT-3′), Sik1-Rv (5′-ACAGG GAGCA AGCAG ATAGG-3′), Sik2-Fw (5′-ACCCC CTCCC TTGAG AGTGT-3′) and Sik2-Rv (5′-CAGACG AGTGG CTCTT GGTGG-3′). Sik1 and Sik2 cDNAs were inserted into Syntegra GEM-T Easy (Promega) and used for digoxigenin (DIG)-labeled probe synthesis. Mice were perfused with PBS followed by 4% paraformaldehyde (PFA), and harvested brains were postfixed in 4% PFA overnight. Forty μm-thick brain sections were treated with 0.3% Triton X-100, digested with 1 μg/ml proteinase K, treated with 0.75% glycine, and then treated with 0.25% acetic anhydride in 0.1 M triethanolamine. After overnight incubation with a DIG-labeled probe at 60 °C, the sections were washed and then incubated with alkaline phosphatase-conjugated anti-DIG Fab fragments (Roche). The reactions were visualized with a
Quantitative RT-PCR and digital PCR. RNA was prepared from the mouse brain, liver, BAT and adrenal gland using the RNasy Lipid Tissue Mini Kit (Qiagen) and QIAzol Lysis Reagent (Qiagen). cDNA was synthesized using oligo dT primer and RNAs as a template with the PrimeScript Reverse Transcriptase Kit (TaKaRa). To quantitate SIK mRNA in each tissue, we used the QX200 Droplet Digital PCR system (Bio-Rad). We prepared a mixture containing cDNA, primers, and ddPCR EvaGreen (Bio-Rad), and processed the mixture for the QX200 Droplet Generator to create droplets. Then, we performed digital PCR and read duplicated droplets using a QX200 Reader according to the manufacturer's instructions.

For quantification of the effect of a high-salt diet, cDNAs were subjected to real-time PCR (ViiA7; ThermoFisher) using SYBR Premix Ex Taq (TaKaRa) and gene-specific primers. The expression of SIK1, 2, and 3 in each tissue, was normalized to the glyceraldehyde-3-phosphate dehydrogenase (GAPDH) gene25. The primers used were Gapdh-Fw (5′-AGAAC ATCAT CCGTG CATCC-3′), Gapdh-Rv (5′-CACAT TGGGG TGAAG AACAC-3′), Sik1-Fw (5′-GACGG AGAGC GTCTG ATACC-3′), Sik1-Rv (5′-GACGC AACCC TTTGA TCTTG-3′), Sik2-Fw (5′-TCCAA GACCT TCTGA GCAGT-3′), Sik2-Rv (5′-CAAGC TTTGA TCTTG AGA-3′), Sik3-Fw (5′-CCAGA GCTCT TCAGA GGGAA-3′) and Sik3-Rv (5′-CGTTT GAGTG GAGA CACTG-3′).

High-fat diet and body weight. Body weight was measured weekly from the age of 6 weeks to 20 weeks. At 6 weeks of age, mice were assigned to a chow diet (MF; Oriental Yeast) or a high-fat diet (D12492; Research Diets). The normal chow diet (MF; Oriental Yeast) provided 3.6 kcal/g (61% carbohydrate, 26% protein, and 13% fat), and the high-fat diet provided 5.2 kcal/g (20% carbohydrate, 20% protein, and 60% fat).

Circadian behavior. Male mice were housed individually in a cage containing a wireless running wheel (Med Associate #ENV-047). The rotation numbers of the wheels were collected by Wheel Manager software (Med Associate) via a receiving device (Med Associate #DIG807). The mice were entrained on the running wheels on a 12 h: 12 h light-dark cycle for 7 days and then allowed to run freely for 14 days under constant darkness. For analysis of re-entrainment to a new light-dark cycle, the mice were maintained on a 12 h: 12 h light-dark cycle, and then the cycle was advanced by six hours. Ten days later, the cycle was advanced again by six hours. The circadian period and re-entrainment were analyzed by a Python-based program. The onset of activity on each day was used to measure the phase relative to the light-dark cycle26. The free-running period was calculated with linear regression analysis of activity onset using a Python-based program. The circadian activity amplitude was normalized to the mean amplitude of the wild-type group4.

Cell culture and plasmids for transfection. HEK293 cells (RC2202) were obtained from the RIKEN BRC Cell Bank. HEK293 cells were cultured and passaged under 5% CO2 in DMEM containing 10% fetal bovine serum and penicillin/streptomycin. Then, 3xFLAG-tagged mouse SIK1, mouse SIK2, and mouse SIK3 cDNAs were cloned into the pcDNA3.1 vector using an In-Fusion HD Cloning Kit (TaKaRa). Although there are multiple protein isoforms of SIK3, we found that SIK3 isoform ending at exon 14 (e.g. UniProt: Q6P4S6-2) is a major form (Ikkkyu et al., unpublished data). We used this SIK3 isoform in this study. Expression vectors for FLAG-SIK1 (S577A) and FLAG-SIK2 (S587A) were generated from pcDNA3-FLAG-SIK1 and pcDNA3-FLAG-SIK2 using a KOD-Plus-Mutagenesis Kit (Toyobo) and a PrimeSTAR Mutagenesis Basal Kit (TaKaRa). Cells were grown to 80% confluency in 6-well plates. HEK293 cells were transfected with 3 μg of pcDNA for FLAG-SIK1, FLAG-SIK2 S577A, FLAG-SIK2 S587A, 3xFLAG-SIK3, and 3xFLAG-SIK3 S551A by using 9 μl of FuGENE HD Transfection Regent (Promega).

Immunoprecipitation and western blot. Used primary antibodies were anti-DYKDDDDD tag antibody (mouse, Monoclonal, Clone #1I6E, Wako) as anti-FLAG antibody, anti-phospho-PKA substrate (RRXS/T) antibody (rabbit, monoclonal, 100G7E, Cell Signaling Technology #9624), anti-14-3-3 (pan) antibody (rabbit, monoclonal, Cell Signaling Technology #5174) and anti-SIK2 antibody (rabbit, polyclonal, Cell Signaling Technology #6919 and Merck #07-1378). Since the anti-SIK2 antibody (Cell Signaling Technology #6919) showed a band similar in size to SIK2 in brains, which was found for both the wild-type and SIK2 KO mice (Fig. S4b), we used the anti-SIK2 antibody (Merck #07-1378) for densitometric analysis of the brain and BAT (Fig. S4c,d, S1). Mice were sacrificed by cervical dislocation under deep anesthesia with sodium pentobarbital (50 mg/kg body weight). Then, BAT and brains were rapidly removed, frozen in liquid nitrogen and stored at −80 °C until use. The BAT and brain were homogenized using a rotor-stator homogenizer (Polytron) in ice-cold lysis buffer (20 mM HEPES pH 7.5, 100 mM NaCl, 10 mM Na2HPO4, 1.5% Triton X 100, 15 mM NaF, 1X PhosSTOP (Roche), 5 mM EDTA, 1X Protease Inhibitor (Roche)), and then centrifuged for 12 min at 14,000 g at 4 °C. For detection of SIK2, brain homogenates were processed for methanol/chloroform precipitation. The pellets were resolved in SDS-PAGE sample buffer. For immunoprecipitation (IP), transiently expressed 3xFLAG-tagged SIK1, SIK2, and SIK3 proteins were extracted with lysis buffer and then centrifuged for 12 min at 14,000 g at 4 °C. The extracted proteins were incubated with a precipitating antibody at 4 °C for 2 h. The beads were washed three times with IP buffer, and the final precipitates were subjected to immunoblotting analysis. The supernatants were separated by SDS–PAGE and transferred to PVDF membranes. The blots were subsequently washed in PBS plus 0.1% Tween-20 (PBS-T) and incubated overnight at 4 °C with primary antibody (1:1000) in PBS-T with 5% bovine serum albumin. The blots were then washed and incubated with horseradish peroxidase-conjugated donkey anti-rabbit IgG (Jackson Immuno Research; 1:5,000).
Statistics. Statistical analyses were performed using SPSS Statistics 22 software (IBM) and Prism 8.0 (GraphPad). All data were subjected to the D’Agostino-Pearson (Omnibus K2) normality test for Gaussian distribution and variance. We performed one-way ANOVA followed by Tukey’s test, one-way repeated measures ANOVA, two-way ANOVA, and two-tailed t-test with Bonferroni correction. We performed one-way repeated measures ANOVA for hourly analysis. P < 0.05 was considered statistically significant. Data were presented as mean ± SEM.

Data availability All data are available from the corresponding author or available on the Open Science Framework (https://osf.io/upgxm/).

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Competing interests
The authors declare no competing interests.

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