mTOR signaling in VIP neurons regulates circadian clock synchrony and olfaction

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Mammalian/mechanistic target of rapamycin (mTOR) signaling controls cell growth, proliferation, and metabolism in dividing cells. Less is known regarding its function in postmitotic neurons in the adult brain. Here we created a conditional mTOR knockout mouse model to address this question. Using the Cre-LoxP system, the mTOR gene was specifically knocked out in cells expressing Vip (vasoactive intestinal peptide), which represent a major population of interneurons widely distributed in the neocortex, suprachiasmatic nucleus (SCN), olfactory bulb (OB), and other brain regions. Using a combination of biochemical, behavioral, and imaging approaches, we found that mice lacking mTOR in VIP neurons displayed erratic circadian behavior and weakened synchronization among cells in the SCN, the master circadian pacemaker in mammals. Furthermore, we have discovered a critical role for mTOR signaling in mediating olfaction. Odor stimulated mTOR activation in the OB, anterior olfactory nucleus, as well as piriform cortex. Odor-evoked c-Fos responses along the olfactory pathway were abolished in mice lacking mTOR in VIP neurons, which is consistent with reduced olfactory sensitivity in these animals. Together, these results demonstrate that mTOR is a key regulator of SCN circadian clock synchrony and olfaction.

mTOR | VIP | SCN | circadian clock | olfaction

Almost all aspects of neuronal functions are regulated by external signals via intracellular signal transduction cascades. Mammalian/mechanistic target of rapamycin (mTOR) is an evolutionarily conserved serine/threonine protein kinase. Centered on mTOR, an intracellular signaling network controls cell growth, proliferation, and metabolism in dividing cells (1, 2). mTOR forms two multiprotein complexes, mTOR complex 1 (mTORC) 1 and mTORC2. mTORC1 activates ribosomal protein S6 kinase (S6K) 1 and S6K2, which in turn phosphorylate the ribosomal protein S6 at Ser240/244 (3–5). mTOR signaling senses intracellular signals including nutrient availability, energy status, and stress, as well as responds to extracellular stimuli by hormones and growth factors. In the developing brain, mTOR signaling promotes neuronal progenitor proliferation, differentiation, and neural circuit formation (6). It is essential in early development, and homozygous mTOR knockout is embryonically lethal in mice (7, 8).

Due to a lack of genetic mouse models of the mTOR mutant, less is known regarding mTOR functions in postmitotic neurons in the adult brain. Studies of mTOR functions were performed using mutants of individual components within mTOR signaling or with pharmacological mTOR inhibitors. It is found that mTOR signaling controls synaptic plasticity, learning, and memory through its interaction with FKBP12 (FK506-binding protein), the mTORC1 downstream effector S6ks, eukaryotic translation initiation factor 4E (eIF4E)-binding protein (4E-BP), and mTORC2 (9–12). mTOR signaling serves as a fuel sensor in the hypothalamus to regulate food intake (13). mTOR also modulates cortical plasticity during sleep and is involved in the effect of sleep deprivation on memory impairment (14, 15). Dysregulation of mTOR signaling pathways in the brain has frequently been identified in neurological and psychiatric disorders (6, 16).

Our previous study pointed to a role for mTOR in the hypothalamic suprachiasmatic nucleus (SCN), the master circadian pacemaker in mammals. The activities of mTORC1 in the SCN exhibit autonomous daily oscillations and are activated by light at night (17, 18). Inhibition of mTOR activity by the drug rapamycin modulates photic resetting of mouse circadian behavior (19). More recently, we have found that mTORC1 promotes mRNA translation of Vip (vasoactive intestinal peptide) via the translation repressor 4E-BP1 (20). VIP is a neuropeptide essential for coupling and synchronization of SCN neurons (21). To further study the functions of mTOR in the SCN as well as in other brain regions, we created a conditional mTOR knockout mouse using the Cre-LoxP system (22). Mtor\textsuperscript{flx/flx} mice (20) were crossed to Vip-Cre mice (23) to specifically knock out mTOR in VIP cells. Using this model, we studied the functions of mTOR in the adult SCN and olfactory bulb (OB), two representative brain regions where VIP neurons are enriched. Using a combination of biochemical, behavioral, and imaging approaches, we demonstrate that mTOR signaling plays a critical role in regulating SCN cell synchrony and olfaction. These results reveal physiological functions of mTOR in the adult brain.

Significance

The mammalian/mechanistic target of rapamycin (mTOR) kinase resides at the crux of an intracellular signaling network that controls fundamental biological processes. Dysregulation of mTOR signaling is linked to neurological and psychiatric diseases. However, the physiological functions of mTOR signaling in the adult brain are not fully understood. In the current study, we discovered that mTOR in vasoactive intestinal peptide (VIP) neurons plays a key role in regulating neurophysiology in the brain circadian clock and the olfactory system. The conditional mTOR knockout mouse will be a useful model for future investigations of mTOR and/or VIP.

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Results

mTOR Is Knocked Down in VIP Neurons. To study the specific role of mTOR in VIP neurons, we crossed Mtorflx/flx mice to Vip-Cre mice to get Mtorflx/flx;Vip-Cre mice. These animals developed normally, were fertile, and did not display gross abnormalities or reduced productivity. Nissl staining indicated that the histological morphology was normal, and the numbers of cells were not decreased in the examined brain regions including the SCN, OB, and piriform cortex in Mtorflx/flx;Vip-Cre mice compared with Mtorflk/flk littermates (Fig. S1). As VIP neurons are enriched in the SCN (24), we first examined mTOR activities in this region by double immunolabeling of VIP and phosphorylated S6 (at Ser240/244; p-S6), a sensitive and specific marker of mTOR activities. We found that p-S6 was strongly expressed in the VIP-expressing cells as well as non-VIP cells in the SCN of Mtorflx/flx mice (Fig. L4). In contrast, p-S6 expression was decreased in the ventral SCN of Mtorflx/flx;Vip-Cre mice, where VIP neurons are located. As a result, the number of cells with colocalized expression of p-S6 and VIP was significantly decreased in the SCN of Mtorflx/flx;Vip-Cre mice, indicating effective knockdown of mTOR activities in VIP neurons. The down-regulation of mTOR was specific, as the number of p-S6–positive non-VIP cells was not changed (Fig. 1 A and B). Next, by Western blotting, we found that mTOR and p-S6 levels were both decreased in the forebrain of Mtorflx/flx;Vip-Cre mice. The level of prepro-VIP, the precursor protein of VIP, was also markedly reduced in the Mtorflx/flx;Vip-Cre mice, whereas the level of the VIP receptor VPAC2 was not changed (Fig. 1 C and D). Together, these results demonstrate that mTOR was specifically knocked down in VIP neurons in the Mtorflx/flx;Vip-Cre mice, and that as a result of mTOR knockdown the prepro-VIP level was also reduced, consistent with a role for mTOR in promoting mRNA translation of Vip (20).

Mtorflx/flx;Vip-Cre Mice Exhibit Abnormal Circadian Behavior. To investigate the functions of mTOR in VIP neurons in the SCN, we first characterized the behavioral phenotypes of the Mtorflx/flx;Vip-Cre mice by recording their circadian wheel-running locomotor activities. Mice were entrained in a 12-h/12-h light/dark (LD) cycle for 7 d and released into constant darkness (DD) for 10 d (Fig. 24). The Mtorflx/flx;Vip-Cre mice were able to entrain to the 12-h/12-h LD cycle and exhibited typical free-running activity rhythms in DD. However, the circadian period in these rhythms was significantly shortened compared with Mtorflk/flk mice (Mtorflk/flk;Vip-Cre vs. Mtorflk/flk: 23.81 ± 0.03 h, n = 20 vs. 23.93 ± 0.02 h, n = 20, t = 3.492, P = 0.0025). After the animals were entrained in LD for 10 d, the LD cycle was abruptly advanced by 8 h. Both groups of mice were able to be reentrained to the shifted LD cycle, but the Mtorflx/flx;Vip-Cre mice were reentrained more quickly than the Mtorflk/flk mice (Mtorflk/flk;Vip-Cre vs. Mtorflk/flk: 7.2 ± 0.35 d, n = 20 vs. 9.45 ± 0.49 d, n = 20, t = 3.319, P = 0.0042).

Next, the mice were released into constant light (LL) for 38 d. Prolonged exposure to LL lengthens the period of circadian behavior and induces arrhythmic behavior in some animals (25). We analyzed the behavioral rhythmicity of the mice using average Fourier periodograms. The average peak of the periodogram of the Mtorflx/flx;Vip-Cre mice was lower than that of the Mtorflk/flk mice, indicating weakened rhythmicity in the Mtorflx/flx;Vip-Cre mice.

Fig. 1. mTOR is knocked down in VIP neurons in the SCN of Mtorflx/flx;Vip-Cre mice. (A) Confocal microscopic images of immunofluorescent labeling for p-S6 (red) and VIP (green) in the suprachiasmatic nucleus. For these experiments, the mice were entrained to a 12-h/12-h light/dark cycle and killed at ZT6. p-S6 and VIP expression was colocalized in the ventral SCN in Mtorflx/flx mice. Note that the number of cells coexpressing p-S6 and VIP (yellow) was decreased in the SCN of Mtorflx/flx;Vip-Cre mice. (B) Representative Western blots of forebrain lysates. (C) Quantitation of the blot intensities is shown. Values are presented as the mean ± SEM. Note that the level of prepro-VIP was markedly reduced in the Mtorflx/flx;Vip-Cre brain but VPAC2 level was not changed. mTOR and p-S6 levels were also significantly reduced. Four Mtorflx/flx and four Mtorflx/flx;Vip-Cre mice were used in the experiment. *P < 0.05 vs. Mtorflx/flx.
(Fig. 2B). We also analyzed the rhythmicity using a discrete wavelet transform, which decomposes the activities into circadian and ultradian components. We used the following two measures (26): (i) cycle-to-cycle variability in period (SD in time between peaks for the circadian component of activity), an indicator of rhythm stability; and (ii) proportion of variance in the activity time series accounted for by the circadian component, which indicates how well-consolidated activity is as a circadian rhythm. A permutation test showed that the cycle-to-cycle variability was significantly higher in the Mtorflx/flx;Vip-Cre mice than in the Mtorflx/mice (P = 0.006; Fig. 2C), indicating less stability of rhythmic behavior in Mtorflx/flx;Vip-Cre mice. The proportion of variance accounted for by the circadian component was lower in the Mtorflx/mice than the Mtorflx/flx;Vip-Cre mice (P = 0.055; Fig. 2D), indicating more spread activities in LL in Mtorflx/flx;Vip-Cre mice. Together, these results indicate more unstable and weakened rhythmicity of Mtorflx/flx;Vip-Cre mice in LL.

**Fig. 2.** Altered circadian behavior in Mtorflx/flx;Vip-Cre mice. (A) Representative double-plotted actograms of mouse wheel-running activities from one Mtorflx/flx (Left) and one Mtorflx/flx;Vip-Cre (Right) mouse. The x axis (Top) indicates the ZT of the day. The y axis (Left) indicates the number of days during the experiment. For these experiments, mice were first entrained to 12-h/12-h light/dark cycles for 7 d and then released into constant darkness for 10 d. Next, animals were reentrained to 12-h/12-h LD for 10 d, and then the LD cycle was abruptly advanced by 8 h. Twenty days later, the LD cycle was delayed by 8 h. After 11 d in the delayed LD cycle, the mice were released into constant light for 38 d. Yellow areas indicate light periods. (B) Averaged Fourier periodograms in LL from all mice. Fourteen Mtorflx/flx and 14 Mtorflx/flx;Vip-Cre mice were used in the experiment. Note that the overall rhythmicity of the Mtorflx/flx;Vip-Cre mice was weaker compared with the Mtorflx/flx mice, as indicated by a lower mean power spectral density (normalized to show proportion power at each frequency). Whereas Mtorflx/mice demonstrated an average of 77.9% of their total activity at the time corresponding to night in the 12-h/12-h LD cycle, Mtorflx/flx;Vip-Cre mice restricted only 40.6% of their total activity to this period (Fig. 2 E and F). These results suggest that multiple oscillatory components exist in the clock of Mtorflx/flx;Vip-Cre mice, one of which cannot be entrained by the 1-h/11-h/1-h/11-h LDLD cycle.
Circadian Synchrony Is Disrupted in the SCN of Mtorflx/flx::Vip-Cre Mice. VIP mediates circadian synchrony among SCN neurons (21). To investigate the cellular basis underlying the abnormal circadian behavior in the Mtorflx/flx::Vip-Cre mice, we crossed these mice to PER2::LUCIFERASE (PER2::LUC) mice (29). In PER2::LUC mice, the luciferase gene is fused to the endogenous clock gene Per2 to create a real-time reporter of dynamic circadian PER2 expression. As previously described (26), we performed microscopic bioluminescence imaging using SCN slices from the PER2::LUC::Mtorflx/flx::Vip-Cre and PER2::LUC::Mtorflx/flx mice. The Mtorflx/flx SCN exhibited pronounced circadian cycles of PER2-driven bioluminescence from abundant bioluminescent regions of interest (ROIs; indicating rhythmic cells) (159 ± 70 ROIs per SCN from four slices). The cellular bioluminescence rhythms were highly synchronized from different ROIs, evident from raster plots (Fig. 3A, Left). In contrast, Mtorflx/flx::Vip-Cre slices exhibited lower-intensity and fewer rhythmic ROIs across the SCN (106 ± 24 ROIs per SCN from six slices), which is consistent with decreased PER1 and PER2 levels in the SCN as determined by immunostaining (Fig. S2). In contrast to the highly synchronized bioluminescence rhythms of ROIs in the Mtorflx/flx slices, ROIs from the Mtorflx/flx::Vip-Cre SCN were less well synchronized, especially during later cycles (Fig. 3A, Right).

Next, we calculated the synchronization (sync) index for each slice, which quantifies the degree of phase clustering among cells, ranging from 0 (uniformly distributed phases across the day) to 1.0 (all cells peak at the same time of day) (30). We found that the sync index was significantly decreased in the SCN of Mtorflx/flx::Vip-Cre mice (Fig. 3C).

To complement the genetic approach to knocking down mTOR, we examined the effects of an mTOR inhibitor, PP242, on SCN cell synchrony. PP242 is a potent, selective, and ATP-competitive mTOR inhibitor; 1 μM PP242 has been shown to significantly inhibit mTOR activities and perturb the translatome in cells (24). In contrast to cellular bioluminescence from slices treated with DMSO (CTR), rhythmic ROIs in PP242-treated SCN were less synchronized, with phases drifting apart over time (Fig. 3B). The sync index was significantly lower in PP242-treated SCN compared with controls (Fig. 3D). Moreover, the proportion of rhythmic ROIs (indicating rhythmic cells) was markedly lower in PP242-treated SCN explants (Fig. 3E), indicating that mTOR is also critical for cellular rhythmicity in the SCN. Together, these results demonstrate that mTOR controls synchrony of SCN cells at least partially through VIP neurons.

Odor Stimulates mTOR Activation in the Olfactory System. The olfactory bulb (OB) is another representative brain region where VIP neurons are enriched (23, 31, 32). Interestingly, VIP has been shown to mediate circadian rhythms in the OB (33). This prompted us to investigate the potential role for mTOR in olfaction using the Mtorflx/flx::Vip-Cre mice.

Fig. 3. mTOR inhibition desynchronizes SCN neurons. (A) Representative raster plots show the daily expression pattern of PER2::LUCIFERASE from 25 representative ROIs (indicators of rhythmic cells) in an SCN slice from an Mtorflx/flx (Left) or Mtorflx/flx::Vip-Cre (Right) mouse. Note that the cellular rhythms of Mtorflx/flx::Vip-Cre were more loosely synchronized compared with the Mtorflx/flx mice. (B) Representative raster plots show the daily expression pattern of PER2::LUCIFERASE from 25 representative ROIs in an SCN slice treated with DMSO (CTR) or the mTOR inhibitor PP242 (1 μM). Note that PP242 significantly disrupted synchronization of SCN cells. (C and D) Quantification of cellular synchronization within each slice using the synchronization index. (E) Percentages of rhythmic ROIs in DMSO (CTR) or PP242-treated SCN slices. Data in C–E are presented as individual values as well as mean ± SEM. For C and D, four Mtorflx/flx SCN slices and six Mtorflx/flx::Vip-Cre slices were used in the experiment. For E, seven SCN slices were used for control and six slices were used for PP242 treatment.
Various odors reach respective olfactory receptors on the olfactory receptor neurons, which project axons to separate glomeruli in the OB and synapse on mitral cells, the principal output neurons of the OB. Olfactory information from the OB is relayed to pyramidal cells in the piriform cortex (PIR). The anterior olfactory nucleus (AON) is also an olfactory relay station which has reciprocal connections with both the OB and PIR. We first examined the expression pattern of p-S6 and its regulation in these regions of Mtor<sup>flx/flx</sup> mice. p-S6 was expressed throughout all layers in the OB, including the periglomerular layer (PGL), external plexiform layer (EPL), mitral cell layer (MCL), and granule cell layer (GCL), as well as in the AON and PIR (Fig. S3). In the OB, p-S6 was enriched in the PGL, MCL, and GCL (Fig. S3A). To investigate potential regulation of mTOR activity by neuronal activities, mice were exposed to an odorant (eucalyptus essential oil) for 15 min at circadian time (CT)15 and killed 45 min after odor exposure. Immunostaining for p-S6 demonstrates that odor evoked significant up-regulation of S6 phosphorylation in the OB, AON, as well as PIR (Fig. S3 A, B, and D). Western blotting confirmed the immunostaining results that odor evoked p-S6 up-regulation in the OB (Fig. S3C).

As reported previously, VIP is extensively expressed in the PGL and EPL and sparsely in the GCL of the OB (31–33). Colocalization of VIP and p-S6 expression was seen in cells in the PGL and EPL (Fig. 4A). In Mtor<sup>flx/flx</sup>;Vip-Cre mice, p-S6 expression was significantly reduced in the VIP-expressing EPL but not in the non–VIP-expressing MCL (Fig. 4B), indicating specific knockdown of mTOR in VIP neurons. Consistent with this, odor-evoked p-S6 up-regulation was abolished in the EPL of Mtor<sup>flx/flx</sup>;Vip-Cre mice (Fig. 4C).

**Olfaction Is Impaired in Mtor<sup>flx/flx</sup>;Vip-Cre Mice.** c-Fos is a marker of neuronal activation and has been shown to be induced by odor in the rodent olfactory system (34–37). To evaluate whether mTOR in VIP neurons mediates olfactory responses, we examined odor-induced c-Fos expression in Mtor<sup>flx/flx</sup> and Mtor<sup>flx/flx</sup>;Vip-Cre mice. As previously reported (37), odor induced c-Fos expression in the OB, AON, and PIR (Fig. 5 A–C). Strikingly, c-Fos induction by odor was abolished in these regions in the Mtor<sup>flx/flx</sup>;Vip-Cre mice (Fig. 5 C and E), indicating an essential role for mTOR signaling in mediating odor-stimulated neuronal responses in the olfactory system. To access the olfactory function of these mice, olfactory sensitivity to a neutral odor (cinnamon) was tested using a behavioral approach as previously described (38). Interestingly, Mtor<sup>flx/flx</sup>;Vip-Cre mice exhibited decreased exploration time toward the odor compared with Mtor<sup>flx/flx</sup> littermates, which indicates decreased sensitivity to the odor. The minimum odor concentration to cause lengthened exploration was increased from 1:1,000 for Mtor<sup>flx/flx</sup> mice to 1:10 for Mtor<sup>flx/flx</sup>;Vip-Cre mice (Fig. 5F).

**Discussion**

To study the physiological function of mTOR in the adult brain, we established a mouse model in which mTOR is specifically knocked down in VIP neurons. Using this model, we found that mTOR signaling in VIP neurons is critical for SCN cell synchronization and olfaction. Mice with decreased mTOR activities in VIP neurons show impaired circadian behavior, which can be mechanistically explained by weakened synchronization among SCN cells. These results are consistent with a role for mTOR in promoting VIP mRNA translation and decreased VIP levels in mTOR knockout animals. Moreover, we found a key role for mTOR in mediating olfactory perception. mTOR activities in the olfactory system are stimulated by odor. mTOR knockout in VIP neurons abolishes odor-evoked c-Fos expression in olfactory regions including the OB, AON, and PIR, which is consistent with reduced olfactory sensitivity in these animals. Together, these results highlight diverse physiological functions of mTOR in the adult brain.

![Fig. 4](image-url)
mTOR is ubiquitously expressed in the brain. mTOR signaling is increasingly found to be involved in unique neuronal activities. As constitutive mTOR knockout is embryonically lethal, studies of mTOR functions in the brain rely on conditional mTOR knockout models. The Cre-LoxP system is commonly used to circumvent embryonic lethality caused by systemic inactivation of a specific gene, since deletion of the gene occurs only in cells where Cre recombinase is expressed. It provides the best experimental control linking genotypes to phenotypes. In our study, Mtor<sup>flx/flx</sup>:Vip-Cre mice were compared with Mtor<sup>flx/flx</sup> littermates. Of note, Vip-Cre is only expressed at the late embryonic to neonatal period, and thus it can be used to manipulate VIP neurons without disruption of their neural development (23). Indeed, the Mtor<sup>flx/flx</sup>:Vip-Cre mice survived without visible deficits, and their brain histology is largely normal.

VIP is a peptide of 28 amino acid residues that belongs to a glucagon/secretin superfamily, the ligand of G protein-coupled receptors. Vip is expressed in a subset of GABAergic neurons in the neocortex, SCN, OB, and other midbrain and brainstem regions as well as the gut and pancreas (39–41). The direct protein product of the Vip gene is prepro-VIP, a 170-amino acid peptide. Our previous work discovered that mRNA translation of Vip is dependent on eIF4E and inhibited by 4E-BPs (20). Binding of 4E-BPs to eIF4E causes inhibition of cap-dependent translation initiation and is relieved when 4E-BPs are phosphorylated by mTOR. Thus, mTOR signaling promotes Vip mRNA translation (20). Knockdown of mTOR in VIP cells markedly decreased the level of prepro-VIP and VIP. Consistently, the mice exhibited significant circadian phenotypes, which largely resemble those seen in Vip or VPAC2 null mice (27,
28, 42, 43) as well as in rats treated with VIP antagonists (44). These phenotypes are (i) weakened circadian rhythmicity under constant conditions; (ii) disrupted circadian behavior under the skeleton photoperiod; and (iii) decreased synchrony among SCN cells. However, as a residual amount of VIP is still expressed in Mtor<sup>flx/flx</sup>:Vip-Cre mice (possibly due to other translational control mechanisms and/or the mosaic pattern of Vip-Cre expression), the VIP-related circadian phenotypes are not as strong as those in Vip and VpAC2 null mice.

Interestingly, the mTOR inhibitor PP242 has a similar, if not stronger, effect on SCN cell synchrony as VIP neuron-specific knockdown of mTOR. As mTOR inhibition by PP242 leads to decreased VIP expression (20), the effect of PP242 may be due to a decreased VIP level in the SCN. However, other mechanisms whereby mTOR regulates SCN cell synchrony cannot be excluded. For example, mTOR inhibition by rapamycin increases GABAergic synaptic transmission (45). As endogenous GABA has been shown to desynchronize circadian cells in the SCN (46), mTOR may also regulate synchrony through modifying GABAergic neurotransmission in the SCN.

mTOR regulates the circadian clock through complex mechanisms. In the current study, genetic and pharmacological inhibition of mTOR activity significantly reduced cellular synchronization as well as the number of rhythmic cells in the SCN. We previously demonstrated that rapamycin inhibits light-induced clock protein PER1 and PER2 expression in the SCN (19). A recent study pointed to the role for mTOR in regulating phototransduction of the clock protein BMAL1 (47). On the other hand, mTOR activity is rhythmically controlled by the circadian clock in the SCN. mTOR activity is high during the day and low at night, which is consistent with close cellular colocalized expression of p-S6 and PER1 in the SCN (18). Light at night rapidly activates mTOR signaling in the SCN (17). Thus, the circadian clock controls the rhythmicity and activity of mTOR. In turn, mTOR signaling feeds back to the circadian clock to regulate its entrainment and synchronization. As mTOR forms a complex signaling network, further in vitro and in vivo studies using specific mutants of mTOR targets will be required to delineate the complex mechanisms whereby mTOR signaling interacts with the circadian clock.

In the current study, the function of mTOR was studied in the olfactory system. We found strong mTOR activities in all layers of the OB as well as in the AON and PIR. Moreover, mTOR activities are stimulated by odorants, and mTOR knockdown in VIP cells diminishes odor-evoked c-Fos responses in the olfactory system. The mechanisms whereby mTOR regulates olfactory c-Fos expression are not clear. A prominent feature of VIP interneurons is their preferential innervation of other interneurons (48, 49). VIP-containing cells in the EPL form an interneuronal network that modulates the function of other inhibitory interneurons (33, 50). VIP neurons may be involved in regulating network excitability and disinhibition of principal cells in the OB, which is similar to a role for VIP neurons in the cortex (51, 52). Thus, the EPL might serve as the “amplifier” of the incoming olfactory signal. mTOR signaling regulates neuronal excitability in cultured neurons (53, 54) as well as in epilepsy animal models (55). mTOR may regulate the excitability of VIP interneurons and therefore control olfactory input through the VIP interneuron network in the EPL. Further electrophysiological studies are required to test these hypotheses.

The regulation and functions of mTOR activity exhibit great similarities between the SCN and OB. First, the base level of mTOR activities is high in both regions. In the SCN, p-S6 expression shows circadian rhythmicity, with the peak level at CT12. Also, the SCN is the only brain region that exhibits a high level of phospho-4E-BPs, other downstream effectors of mTORC1. Second, mTOR activities are induced by neuronal activities in both regions. In the SCN, VIP cells are among the photic recipient neurons that receive direct synaptic input from the intrinsically photosensitive retinal ganglion cells. mTOR signaling is rapidly activated in these neurons by light stimulation at night. Similarly, odor stimulation activates mTOR in neurons in the OB. Third, mTOR exhibits similar functions in the SCN and OB in its regulation of neuronal network properties. In the SCN, VIP neurons are in the ventral region, but they form a neuronal network that couples and synchronizes the ventral and dorsal regions of the SCN. In the OB, VIP-expressing interneurons form a network in the EPL that is critical for controlling olfactory input. Thus, mTOR signaling regulates properties of structured VIP neuronal networks in the SCN and OB.

### Materials and Methods

#### Animals

Mtor<sup>flx/flx</sup>:Vip-Cre mice on a C57BL/6 background kindly provided by Sara C. Kozma, University of Cincinnati, Cincinnati, OH, were crossed to a VIP promoter-driven Cre-recombinase mouse line (Vip-Cre; Jackson Laboratory) to generate Mtor<sup>flx/flx</sup>:Vip-Cre mice. Mice were then crossed with mPER2::LUC transgenic reporter mice (29) to obtain Mtor<sup>flx/flx</sup>:Vip-Cre:mPER2::LUC mice. Mice were maintained in the animal facilities at the University of Minnesota, Duluth Campus, McGill University, or Concordia University in accordance with institutional guidelines. All procedures were approved by the Institutional Animal Care and Use Committees at University of Minnesota, McGill University, and Concordia University.

#### Brain Tissue Processing, Immunostaining, and Microscopic Imaging Analysis

Under the indicated conditions, the mice were killed and brain tissue was harvested. Brain sections were processed and immunostained for p-S6, VIP, and c-Fos as previously reported (45). Bright-field and fluorescent microscopic images were captured using a digital camera mounted on an inverted DMi8 Leica microscope. Confocal microscopy images were captured using a Zeiss 710 Meta confocal microscope. See Table 1 for antibody information.

All photomicrographic datasets were statistically analyzed using Adobe Photoshop software (Adobe Systems). For the p-S6 and VIP colocalization assay, confocal SCN images (40x magnification) of double labeling for p-S6 and VIP were collected. Individual SCN cells were outlined based on DRAQ5 (a cell nuclear dye) staining, and the expression of p-S6 (red), VIP (green), or both (yellow) was determined based on densitometry values for red (p-S6)

### Table 1. Antibodies used for immunostaining and Western blotting

| Antibody | Supplier | Catalog no. | Dilution in immunostaining | Dilution in Western blotting |
|----------|----------|-------------|---------------------------|-----------------------------|
| VIP      | Santa Cruz | sc-21041    | 1:200                     |                             |
| mTOR     | Cell Signaling | 2983      | 1:1,000                   |                             |
| p-S6     | Cell Signaling | 2215      | 1:1,000                   |                             |
| S6       | Santa Cruz | 74459       | 1:2,000                   |                             |
| PER1     | MilliporeSigma | A82201   | 1:3,000                   |                             |
| PER2     | Santa Cruz | SC-7728     | 1:300                     |                             |
| Prepro-VIP | Sigma-Aldrich | V0390   | 1:500                     |                             |
| VPAC2    | Abcam     | ab28624     | 1:1,000                   |                             |
| β-Actin  | Sigma-Aldrich | A5441   | 1:5,000                   |                             |
| c-Fos    | Calbiochem | PC38        | 1:3,000                   |                             |

See Table 1 for antibody information.
Six- to 8-wk-old mice were entrained to a 12-h/12-h LD cycle for 10 d and transferred to a skeleton photoperiod, where no p-S6 was expressed, to determine the intensity of non-specific background staining. The background value was subtracted from the p-S6 labeling value to obtain the normalized p-S6 intensity. Three brain sections were used from each animal and three mice were used in each group.

Protein Extraction and Western Blotting Analysis. Total forebrain tissue was homogenized with a pestle grinder (Fisher Scientific) and lysed using a lysis buffer as previously reported (20). Western blotting analysis was performed and the intensity of the blots was analyzed as described (56). See Table 1 for antibody information.

Circadian Behavioral Assay. Eight- to 10-wk-old male mice were individually housed in cages equipped with running wheels. Wheel rotation was recorded using the VitalView program (Mini Mitter) or ClockLab software (Actimetrics) (45). The animals were entrained to a 12-h/12-h light/dark cycle for 9 d, re-synchronized using the VitalView program (Mini Mitter) or ClockLab software (Actimetrics) – previously described (26). Briefly, sections containing SCN (150 μm) were collected, cultured on a membrane (Millicell CM; Millipore) in 1.2 mL of air-buffered media containing 0.1 mM bmeet luciferin (Gold Biotechnology), and imaged for 5 d using a Stanford Photonics XR/MEGA-102 cooled intensified charge-coupled device camera. For pharmacological experiments, the mTOR inhibitor PF242 or DMSO was added to respective culture media and dishes were imaged simultaneously.

Rhythmic parameters of PER2::LUC expression were calculated for each slice and for cell-like regions of interest (ROIs) within each slice using MATLAB (MathWorks)-based computational analyses as described previously (46). Briefly, phase maps of slices were constructed by generating a 4-d time series for each 12-pixel-diameter region of the image that met the criteria for circadian rhythmicity, namely a peak autocorrelation coefficient significant at alpha = 0.05 and associated with lag between 18 and 34 h. To locate and extract data from cell-like ROIs, an iterative process was employed after background and local noise subtraction of a slice image summed across 24 h of bioluminescence (26). The synchronization index R is computed as in equation 6 of ref. 30, using 2-d segments. To assess overall rhythmicity in each slice, 4-d time series were extracted for each of the 20% brightest pixels in each processed image (with bioluminescence summed across 24 h; a total of 5,369 pixels used for each slice) and evaluated for rhythmicity using the same criteria as above. The power density of these pixels associated with significantly rhythmic time series is used as a general estimate of what proportion of the slice is rhythmic. Because of small sample sizes and non-normal distributions, permutation tests were used to test for significant difference in mean between groups. Raster plots show 36 cell-like ROIs chosen by sorting the first peak times and selecting 36 evenly spaced ROIs from the sorted list to obtain a more representative sample.

Olfactory Behavioral Test. Olfactory sensitivity was tested as described previously (38). Briefly, at zeitgeber time (ZT)15, 6- to 8-wk-old mice were exposed to one of four different dilutions of cinnamon extract (Watkins) or water on filter paper for a 3-min session. Total time spent exploring the filter paper was assessed.

Statistical Analysis. Values are presented as the mean ± SEM or percentage. Statistical analysis was performed using SPSS software (SPSS). Mean values from multiple groups were compared via one-way ANOVA, followed by Bonferroni’s multiple comparisons. Mean values from two groups were compared via Student’s t test. P < 0.05 was considered statistically significant.

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Explain Culture, Kinetic Bioluminescence Imaging, and Data Analysis. Explants of SCN tissues from MtorflVip-Cre:mPER2::LUC and MtorflVip-Cre::mPER2::LUC mice were dissected and cultured as reported (26). For cellular-resolution real-time assays, coronal sections containing SCN were imaged as previously described (26). Briefly, sections containing SCN from each mouse were collected, cultured on a membrane (Millicell CM; Millipore) in 1.2 mL of air-buffered media containing 0.1 mM bmeet luciferin (Gold Biotechnology), and imaged for 5 d using a Stanford Photonics XR/MEGA-102 cooled intensified charge-coupled device camera. For pharmacological experiments, the mTOR inhibitor PF242 or DMSO was added to respective culture media and dishes were imaged simultaneously.

Rhythmic parameters of PER2::LUC expression were calculated for each slice and for cell-like regions of interest (ROIs) within each slice using MATLAB (MathWorks)-based computational analyses as described previously (46). Briefly, phase maps of slices were constructed by generating a 4-d time series for each 12-pixel-diameter region of the image that met the criteria for circadian rhythmicity, namely a peak autocorrelation coefficient significant at alpha = 0.05 and associated with lag between 18 and 34 h. To locate and extract data from cell-like ROIs, an iterative process was employed after background and local noise subtraction of a slice image summed across 24 h of bioluminescence (26). The synchronization index R is computed as in equation 6 of ref. 30, using 2-d segments. To assess overall rhythmicity in each slice, 4-d time series were extracted for each of the 20% brightest pixels in each processed image (with bioluminescence summed across 24 h; a total of 5,369 pixels used for each slice) and evaluated for rhythmicity using the same criteria as above. The power density of these pixels associated with significantly rhythmic time series is used as a general estimate of what proportion of the slice is rhythmic. Because of small sample sizes and non-normal distributions, permutation tests were used to test for significant difference in mean between groups. Raster plots show 36 cell-like ROIs chosen by sorting the first peak times and selecting 36 evenly spaced ROIs from the sorted list to obtain a more representative sample.
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