AMP-Activated Protein Kinase Regulates Circadian Rhythm by Affecting CLOCK in *Drosophila*

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The circadian clock organizes the physiology and behavior of organisms to their daily environmental rhythms. The central circadian timekeeping mechanism in eukaryotic cells is the transcriptional-translational feedback loop (TTFL). In the *Drosophila* TTFL, the transcription factors CLOCK (CLK) and CYCLE (CYC) play crucial roles in activating expression of core clock genes and clock-controlled genes. Many signaling pathways converge on the CLK/CYC complex and regulate its activity to fine-tune the cellular oscillator to environmental time cues. We aimed to identify factors that regulate CLK by performing tandem affinity purification combined with mass spectrometry using *Drosophila* S2 cells that stably express HA/FLAG-tagged CLK and V5-tagged CYC. We identified SNF4A γ , a homolog of mammalian AMP-activated protein kinase γ (AMPK γ), as a factor that copurified with HA/FLAG-tagged CLK. The AMPK holoenzyme composed of a catalytic subunit AMPK α and two regulatory subunits, AMPK β and AMPK γ , directly phosphorylated purified CLK *in vitro*. Locomotor behavior analysis in *Drosophila* revealed that knockdown of each AMPK subunit in pacemaker neurons induced arrhythmicity and long periods. Knockdown of AMPK β reduced CLK levels in pacemaker neurons, and thereby reduced pre-mRNA and protein levels of CLK downstream core clock genes, such as *period* and *vri*. Finally, overexpression of CLK reversed the long-period phenotype that resulted from AMPK β knockdown. Thus, we conclude that AMPK, a central regulator of cellular energy metabolism, regulates the *Drosophila* circadian clock by stabilizing CLK and activating CLK/CYC-dependent transcription.

Key words: AMP-activated protein kinase; circadian clock; CLOCK; *Drosophila*; tandem affinity purification; TTFL

Significance Statement

Regulation of the circadian transcription factors CLK and CYC is fundamental to synchronize the core clock with environmental changes. Here, we show that the AMPK γ subunit of AMPK, a central regulator of cellular metabolism, copurifies with the CLK/CYC complex in *Drosophila* S2 cells. Furthermore, the AMPK holoenzyme directly phosphorylates CLK *in vitro*. This study demonstrates that AMPK activity regulates the core clock in *Drosophila* by activating CLK, which enhances circadian transcription. In mammals, AMPK affects the core clock by downregulating circadian repressor proteins. It is intriguing to note that AMPK activity is required for core clock regulation through circadian transcription enhancement, whereas the target of AMPK action is different in *Drosophila* and mammals (positive vs negative element, respectively).

Introduction

The circadian clock allows living organisms to manifest daily rhythms in physiology and behavior by synchronizing internal

processes with rhythmic environmental changes. In eukaryotes, the cell-autonomous circadian clock operates via transcriptional-translational feedback loops (TTFLs) composed of core clock proteins. In *Drosophila*, CLOCK (CLK) and CYCLE (CYC) activate the transcription of *period* (*per*) and *timeless* (*tim*), and PER and TIM proteins, in turn, repress their own transcription in the core loop. CLK and CYC also activate the transcription of *vri* (*vri*) and the PAR domain protein 1 ϵ gene (*Pdp1 ϵ*). VRI and PDP1 ϵ repress and activate the transcription of *Clk*, which enhances the robustness of the interlocked feedback loop. CLOCK-

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WORK ORANGE (CWO), another downstream target of CLK/CYC, works cooperatively with PER to increase the amplitude of oscillation (Hardin, 2011). A similar mechanism operates in mammals with CLK/BMAL1 as the master circadian transcription factors and PER1, PER2, and PER3, and CRYPTOCHROME1 and 2 (CRY1, CRY2) as the transcriptional repressors in the core loop. Nuclear receptors ROR α and REB-erb α , which are induced by CLK/BMAL1, activate and inhibit the expression of BMAL1, respectively, to form an interlocked feedback loop (Partch et al., 2014).

In the *Drosophila melanogaster* brain, ~150 pacemaker neurons strongly express the core clock genes and drive the locomotor activity rhythms (Helfrich-Förster, 2005). The pacemaker neurons form clusters, and each cluster performs a specific function that determines activity profiles in response to the environmental inputs. The dorsal neurons are DN1_s, DN1_ps, DN2s, and DN3s. The lateral clusters are the ventrally located neurons (LN_vs), including the large lateral ventral neurons (LLN_vs), the small lateral ventral neurons (sLN_vs), the fifth LN_v, and the lateral dorsal neurons (LN_ds). Among the pacemaker neurons, the sLN_vs are crucial for rhythmic behavior under constant conditions (Helfrich-Förster, 1998; Renn et al., 1999; Nitabach et al., 2002). Further, sLN_vs project dorsally and terminate in the vicinity of DN1s and DN2s and express peptide dispersing factor (PDF), a neuropeptide that allows sLN_vs to coordinate pacemaker neurons to maintain synchrony (Renn et al., 1999; Lin et al., 2004; Helfrich-Förster et al., 2007).

Because CLK is the rate-limiting component of TTFL, maintaining the appropriate CLK activity and level is crucial for circadian rhythm generation. Increasing the transcription activation potency of CLK increases the speed of the circadian clock (Kadener et al., 2008; Lee et al., 2014), and flies harboring the hypomorphic allele *Clk*^{ar} that has reduced CLK transcription activity do not exhibit circadian rhythms (Allada et al., 2003). The phosphorylation pattern of CLK also has daily rhythms that tightly couple with the rhythms of activity and stability. Early day and late night hyperphosphorylation is primarily driven by the *D. melanogaster* kinase DOUBLETIME (DBT), which phosphorylates PER and is the homolog of the mammalian casein kinase I δ/ϵ , reduces CLK stability, and correlates with reduced transcriptional activity (Kim and Edery, 2006; Yu et al., 2006; Menet et al., 2010). Consistently, dephosphorylation of CLK by protein phosphatase 2A (PP2A) induces stabilization and enhances transcriptional activity (Kim and Edery, 2006; Andreatza et al., 2015). NEMO, another kinase that phosphorylates PER, has also been shown to destabilize CLK in flies (Yu et al., 2011; Szabó et al., 2013), but casein kinase 2 (CK2) is recruited onto CLK during late night but makes CLK stable and inactive (Szabó et al., 2013). During mid-day when CLK retains high transcriptional activity, CLK is present in the medium to hypophosphorylated isoforms. This mid-day specific phosphorylation occurs on the 16 Ser/Thr residues by yet unidentified kinases (Lee et al., 2014) and maintains the appropriate levels and transcriptional activity of CLK, which is crucial to synchronize animal behavior to the temperature cycle.

In this study, we aimed to isolate proteins that interact with CLK/CYC to understand how CLK/CYC activity is regulated. *D. melanogaster* S2 cells that stably expressed FLAG/HA epitope-tagged CLK and V5 epitope-tagged CYC were used for tandem affinity purification (TAP) followed by mass spectrometry (MS) analysis. We identified SNF4A γ , the regulatory subunit of the AMP-activated protein kinase (AMPK) holoenzyme, as a protein that interacts with CLK/CYC. Downregulation of each subunit of

the AMPK holoenzyme in pacemaker cells significantly altered circadian locomotor behavior, indicating that the enzymatic activity of AMPK is required for normal circadian clock function. Moreover, the AMPK holoenzyme directly phosphorylated GST-tagged CLK proteins *in vitro*. Because CLK protein levels were reduced in clock cells and because the ectopic expression of CLK rescued the period-lengthening defect of the AMPK β subunit knockdown, we concluded that the major cellular energy sensor AMPK affects the core clock by stabilizing CLK in *D. melanogaster*.

Materials and Methods

Plasmids. To generate the pMT-*Clk*-HA/FLAG plasmid, the *Clk* open reading frame was amplified from the SK(–)/*Clk* plasmid (Lee et al., 1999) and cloned into a pMT-HA/FLAG expression vector carrying a hygromycin resistance cassette (Kusch et al., 2004). The SNF4A γ open reading frame was PCR-amplified from cDNA that had been reverse-transcribed in a manner that excluded the stop codon from total RNA extracted from fly heads, and the PCR product was subcloned into the pMT-V5-His vector (Invitrogen). The PCR primers used were 5'-ATTGGTACCATGCATGGAATAACACACCT-3' and 5'-ATTTCTAGATTACTAACCAACGCCATTT-3'; the SNF4A γ sequences are underlined. The pMT-*cyc*-V5 and pMT-HA-*Clk* plasmids were described previously (Kim and Edery, 2006).

TAP. *D. melanogaster* S2 cells were cotransfected with 9.5 μ g each of the pMT-*Clk*-HA/FLAG and pMT-*cyc*-V5 plasmids using calcium phosphate (Invitrogen) according to the manufacturer's protocol. Stable transgenic S2 cell lines harboring the pMT-*Clk*-HA/FLAG and pMT-*cyc*-V5 plasmids were selected by culturing in a hygromycin-containing medium. Stable transgenic S2 cells were expanded in a 2 L suspension culture, and CLK-HA/FLAG and CYC-V5 expression was induced by the addition of CuSO₄ to a final concentration of 500 μ M. After a 1 d induction, the cells were harvested, and proteins were extracted in lysis buffer [50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 0.4% (v/v) NP-40, 1.5 mM MgCl₂, 5% glycerol, 1 mM EDTA, 1 mM DTT, 1 mM Na₃VO₄, 2.5 mM NaF, and a freshly added protease inhibitor mixture (Sigma-Aldrich)]. Protein extracts were incubated with 0.6 ml of EZview Red anti-FLAG M2 affinity gel (Sigma-Aldrich, RRID:AB_2616449) overnight at 4°C with rotation. The gel was then washed three times with lysis buffer and eluted using the 3 $\times$ FLAG peptide (1.5 mg/ml). The eluate was diluted with lysis buffer and incubated with EZview Red anti-HA affinity gel (Sigma-Aldrich, RRID:AB_10109562) for 3 h at 4°C with rotation. The gel was then washed three times with lysis buffer and eluted using the 2HA peptide (1 mg/ml). The final eluate was used for MS analysis (DiatechKorea).

Nano-liquid chromatography (LC)-MS/MS analysis. Proteins in the eluate were separated by SDS-PAGE and stained with Coomassie Brilliant Blue. Protein bands were excised from the gel, and the gel pieces were vortexed in 50% acetonitrile (ACN) containing 50 mM NH₄HCO₃ until the stain was completely removed. The gel pieces were then dehydrated in 100% ACN and vacuum-dried for 20 min using a SpeedVac. For in-gel digestion, the disulfide bridges were reduced with 10 mM DTT in 50 mM NH₄HCO₃ for 45 min at 56°C, and then the cysteines were alkylated with 55 mM iodoacetamide in 50 mM NH₄HCO₃ for 30 min in the dark. Finally, each gel piece was treated with 12.5 ng/ μ l sequencing-grade modified trypsin (Promega) in 50 mM NH₄HCO₃ buffer, pH 7.8, at 37°C overnight. Following digestion, tryptic peptides were extracted with 5% formic acid in 50% ACN at room temperature for 20 min. The supernatants were collected and dried using a SpeedVac. The samples were resuspended in 0.1% formic acid and purified and concentrated using C18 ZipTips (Millipore) before MS analysis. The samples were loaded onto a fused silica microcapillary column (12 cm $\times$ 75 μ m) packed with C18 reversed-phase resin (5 μ m, 200 Å). LC was conducted using a linear gradient: a 3%–40% gradient of solvent B (0.1% formic acid in ACN) in solvent A (0.1% formic acid in distilled water) with a flow rate of 250 nl/min for 60 min. The column was directly connected to an LTQ linear ion-trap MS (ThermoFinnigan) equipped with a nano-electrospray ion source. Each full MS scan was followed by five MS/MS scans of the most intense to the fifth-most intense peaks of the full MS scan. The repeat

count of peaks for dynamic exclusion was 1, with a 30 s repeat duration. The dynamic exclusion duration was set to 180 s, and the width of exclusion mass was ± 1.5 Da. The list size of dynamic exclusion was 50. The acquired LC-ESI-MS/MS fragment spectra were searched for in the BioWorksBrowser (version Rev. 3.3.1 SP1, Thermo Fisher Scientific, RRID:SCR_014594) with the SEQUEST search engine against the *D. melanogaster* database at National Center for Biotechnology Information (<http://www.ncbi.nlm.nih.gov/>). The following search conditions were used: trypsin enzyme, up to two missed cleavages permitted, peptide tolerance of ± 2 amu, a mass error of ± 1 amu on the fragment ions, and fixed modifications of carbamidomethylation of cysteines (57 Da) and oxidation of methionines (16 Da).

Fly strains. The following fly strains were used: w^{1118} (BDSC, BL5905), *cyc*⁰¹ (Rutila et al., 1998), *Clk*^{out} (Lee et al., 2014), *Clk*^{irk} (Allada et al., 1998), *dcr2*;tim(UAS)-Gal4 (Kim et al., 2012), UAS-*Clk* (Zhao et al., 2003), pdf-Gal4 (Park et al., 2000), tub-Gal80^{ts};tim(UAS)-Gal4 (Kim et al., 2012), UAS-AMPK α Ri-1 (VDRC1827), UAS-AMPK α Ri-2 (NIG3051R-1), UAS-AMPK β Ri-1 (NIG8057R-2), UAS-AMPK β Ri-2 (NIG8057R-4), UAS-AMPK γ Ri-1 (NIG17299R-3), UAS-AMPK γ Ri-2 (NIG17299R-1), UAS-GFP.nls (BDSC, BL4776), and UAS-mCD8::GFP (BDSC, BL5130). pdf-Gal4 was crossed to UAS-*dicer2*/FM6 to generate *dcr2*;pdf-Gal4. For AMPK subunit knockdown, driver flies [e.g., *dcr2*;actin-Gal4, *dcr2*;tim(UAS)-Gal4, *dcr2*;pdf-Gal4, and tub-Gal80^{ts};tim(UAS)-Gal4] were crossed to UAS-AMPK RNAi flies. UAS-RNAi flies used in this study were in the w^{1118} genetic background. Driver flies were crossed to w^{1118} flies and served as experimental heterozygous controls (marked as actin>*d2*, tim>*d2*, pdf>*d2*, and tim,G80^{ts}>*d2*, respectively). None of the fly strains was backcrossed to w^{1118} flies.

Behavioral analysis. The locomotor activity of individual flies was determined using the Drosophila Activity Monitoring system (Trikinetics). Young male adult flies were exposed to 12 h of light and 12 h of dark (LD) for 4 d and then were maintained in constant darkness (DD) for 7 d at the indicated temperatures. The locomotor data analysis was performed using FaasX software (Fly Activity Analysis Suite for MacOSX), which was generously provided by Francois Rouyer (Centre National de la Recherche Scientifique, France). Periods were calculated for each fly using χ^2 periodogram analysis, and data were pooled to obtain an average value. Power was calculated by quantifying the relative strength of the rhythm during DD. Individual flies with a power ≥ 10 and a width ≥ 2 were considered rhythmic. Actograms represent the double-plotted locomotor activities throughout the experimental period and were acquired using Actogram J software (Schmid et al., 2011).

qRT-PCR. qRT-PCR was performed as described previously (Lee et al., 2016). Total RNA was extracted from fly heads using QIAzol reagent (QIAGEN). Total RNA (1 μ g) was reverse-transcribed using an oligo(dT) 20 primer (for mRNA) or a random hexamer primer (for pre-mRNA) and PrimeScript RTase (TaKaRa). Real-time PCR was performed using Rotor Gene 6000 (QIAGEN) with SYBR Premix Ex Taq (Tli RNaseH Plus, TaKaRa). The following primers were used: *ampk α* forward, 5'-CCGGCAAGTCTCGAAGAT-3'; *ampk α* reverse, 5'-GAGTAAGGCTCTCCATGATGAC-3'; *ampk β* forward, 5'-GTGACCAAC TATGCGGAGAA-3'; *ampk β* reverse, 5'-AGCGGTGTATCCTTGTTG AG-3'; *ampk γ* forward, 5'-ATCGAGGACATACCCGAAGA-3'; *ampk γ* reverse, 5'-GTCTGGTCTGCTGGTTATT-3'; *per* forward, 5'-GTGAGAGCGAGAGCGAGTGT-3'; *per* reverse, 5'-TATGTAAGCTGC CTGCCAA-3'; *vri* forward, 5'-CAAGCAGGATAATCCAGCAA-3'; and *vri* reverse, 5'-ACAGATTTCAGATCAAACGTGGA-3'. Non-cycling mRNA encoding *cbp20* was used to normalize gene expression with the following primers: *cbp20* forward, 5'-GTATAAGAAGACGC-CCTGC-3'; and *cbp20* reverse, 5'-TTCACAAATCTCATGGCCG-3'. The data were analyzed using Rotor Gene 6000 software, and the relative mRNA levels were quantified using the $2^{-\Delta\Delta Ct}$ method in which $\Delta\Delta Ct = [(C_t \text{ target} - C_t \text{ cbp20}) \text{ at ZTx}] - [(C_t \text{ target} - C_t \text{ cbp20}) \text{ at ZT4}]$.

Western blotting and immunoprecipitation. For Western blotting, protein extracts were prepared in HEMG lysis buffer [10 mM HEPES, pH 7.5, 5 mM Tris-HCl, pH 7.5, 50 mM KCl, 10% glycerol, 0.1% Triton X-100, 1 mM EDTA, and 1 mM DTT] with freshly added protease inhibitor mixture (Sigma-Aldrich), 1 mM Na₃VO₄, and 1 mM NaF. To detect dCLK, protein extracts were prepared in RIPA lysis buffer [25 mM Tris-HCl, pH

7.5, 50 mM NaCl, 0.5% sodium deoxycholate, 0.5% NP-40, and 0.1% SDS] and were briefly sonicated (Kim and Edery, 2006). Protein extracts were resolved by SDS-PAGE with the indicated polyacrylamide percentage or on 3%–8% Tris-acetate Criterion gels (for CLK). Primary antibodies were used at the following dilutions: anti-CLK (gp208), 1:2000; anti-V5 (Invitrogen, RRID:AB_159298), 1:2000; anti-HA (Roche Diagnostics, RRID:AB_514505), 1:2000; anti-PER (Rb1), 1:2000; anti-TIM (Rb1), 1:2000; and anti-AMPK α (AbD Serotec, RRID:AB_1604624), 1:2000. Band intensity was quantified using ImageJ software (RRID:SCR_003070).

For immunoprecipitation, protein extracts were prepared in modified-RIPA lysis buffer (50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1% NP-40, and 0.25% sodium deoxycholate) and were briefly sonicated. Then, 3 μ l of anti-V5 (Invitrogen, RRID:AB_159298) or anti-HA (Roche Diagnostics, RRID:AB_514506) was added to the extract depending on the target protein and incubated overnight at 4°C with end-over-end rotation. Normal rabbit IgG (Santa Cruz Biotechnology) was used as a negative control. Finally, 20 μ l of Gamma Bind-Sepharose beads (GE Healthcare) was added, and the mixture was further incubated for 3 h at 4°C with end-over-end rotation. The immune complexes were then eluted from the beads using SDS-PAGE sample buffer.

Immunostaining. Immunostaining was performed as previously described (Lee et al., 2016). Briefly, fly heads were fixed in 4% PFA and washed with 0.5% Triton X-100 (PBT). The fixed heads were dissected, the isolated brains were permeabilized in 1% PBT for 20 min, and then blocked in 0.5% PBT containing 5% horse serum for 30 min. The following primary antibodies were added to the mixtures directly: anti-PDF antibody (C7) (Cyran et al., 2005), diluted 1:200; anti-VRI (gp2) (Glossop et al., 2003), diluted 1:1000; anti-PER (Rb1) (Kim et al., 2012), diluted 1:200; and anti-CLK (gp50) (Houl et al., 2006), diluted 1:200. The brains were washed with PBT and incubated with secondary antibodies in a blocking solution overnight at 4°C. The following secondary antibodies were used at 1:200 dilution: goat anti-rabbit Alexa-488 (Thermo Fisher Scientific, RRID:AB_143165), goat anti-mouse Alexa-555 (Thermo Fisher Scientific, RRID:AB_141780), and goat anti-mouse Alexa-633 (Thermo Fisher Scientific, RRID:AB_2535718). Stained brain samples were washed with PBT, incubated in 0.1 M phosphate buffer containing 50% glycerol for 30 min, and mounted using a mounting medium. Confocal images were obtained using an LSM 710 confocal microscope (Carl Zeiss) and processed using the Zen software (ZEN Digital Imaging for Light Microscopy, Carl Zeiss, RRID:SCR_013672). For signal quantification, the pixel intensity of each cell was determined using ImageJ software. Briefly, the level of background staining was measured in the field surrounding the cluster of clock neurons and subtracted from the pixel intensities determined for the clock neuronal cells. The final intensity was the average of at least five brains for each genotype.

In vitro kinase assay. GST-tagged recombinant CLK proteins were purified from *Escherichia coli* BL21 cells, which were transformed with pGEX4T-1-*dClk*, using the MagneGST protein purification system (Promega). Briefly, *E. coli* pellets were lysed with lysis buffer (MagneGST cell lysis reagent, Promega) containing 1 mg/ml lysozyme (Sigma-Aldrich), RNase-free DNase, and 5 mM DTT added fresh. MagneGST particles were equilibrated with lysis buffer and incubated with the lysates for 3 h at 4°C with rotation. Magnetic particles with bound GST-tagged CLK were collected using a magnetic stand and were then washed with lysis buffer three times. Protein was eluted with a 50 mM Tris-HCl, pH 8.1, buffer containing 50 mM glutathione.

The *in vitro* kinase assay was performed with 1 μ g of purified GST-CLK as the substrate and 50 ng active AMPK holoenzyme (Millipore) in 20 mM HEPES, pH 7.5, 10 mM MgCl₂, 1 mM DTT, 6.25 mM β -glycerophosphate, 1.25 mM EGTA, and 0.125 mM cold ATP (or with 12 μ Ci γ -³²P-ATP for the radioactive assay) in the presence or absence of 0.15 mM AMP at 30°C for 20 min. The samples were resolved by SDS-PAGE and then transferred to a PVDF membrane. The membrane was stained with Ponceau S, and phosphorylation of the substrate was visualized by autoradiography.

Statistical analysis. Statistical analysis was performed using Prism version 7.0 software (GraphPad; RRID:SCR_002798). Significant differences between groups were determined using the Student's *t* test, one-

Figure 1. Confirmation of the interaction between AMPK γ and CLK by reciprocal coimmunoprecipitation. **A**, S2 cells that stably expressed TAP-tagged CLK and V5-tagged CYC were treated with (+) or without (–) 500 μ M CuSO_4 (final concentration). Then, 24 h after induction, protein extracts were analyzed by immunoblotting. The expression of TAP-tagged CLK and V5-tagged CYC induced expression of endogenous PER and TIM proteins as shown on Western blots probed with anti-PER antibody (Rb1) and anti-TIM antibody (Rb1). **B**, **C**, S2 cells were transiently transfected with pMT-HA-*Clk* and pMT-SNF4A γ -V5, and expression of HA-CLK and AMPK γ -V5 was induced by the addition of 500 μ M CuSO_4 . Then, 24 h after induction, protein extracts were analyzed directly by immunoblotting (Input) or subjected to immunoprecipitation (IP) with anti-HA antibody, anti-V5 antibody, or anti-normal IgG antibody as a negative control.

way ANOVA, or two-way ANOVA followed by the Tukey's test or the Dunnett's test as indicated in the figure legends.

Results

TAP analysis identified AMPK γ as a protein that interacts with CLK

To identify proteins that interact with CLK, we performed TAP with extracts from S2 cells that stably expressed HA/FLAG tandemly tagged CLK (TAP-tagged CLK) and V5-tagged CYC. We reasoned that because most CLK, if not all, functions as a heterodimer with CYC, coexpression of CYC with CLK would increase the formation of functionally intact CLK-CYC complexes. To determine whether CLK/CYC activity was affected by the tags, PER and TIM protein levels were measured in stable S2 cells. As reported previously (Saez and Young, 1996), the PER and TIM proteins were not detectable in stable S2 cells without the induction of CLK and CYC (Fig. 1A). However, 24 h after induction, PER and TIM protein expression was enhanced and detectable, indicating that the epitope-tagged CLK and CYC were functional and able to activate transcription (Fig. 1A).

We next purified TAP-tagged CLK and the associated proteins from total protein lysates of S2 cells, and identified the associated proteins by LC-MS/MS. In addition to CLK and CYC, the core clock proteins, including TIM, PER, DBT, and Supernumerary Limb, were among the top-hit identified proteins (data not shown), indicating that TAP successfully recovered the functional CLK/CYC complex. Among the associated proteins that were identified, we focused on SNF4A γ , which encodes the γ regulatory subunit of AMPK and is an ortholog of mammalian *Prkag1*, *Prkag2*, and *Prkag3*. SNF4A γ is henceforth referred to as AMPK γ (Pan and Hardie, 2002). The interaction between CLK and AMPK γ was verified by immunoprecipitation; ectopically expressed HA-tagged CLK and V5-tagged AMPK γ were coimmunoprecipitated from S2 cells in a reciprocal manner (Fig. 1B,C).

Knockdown of expression of each AMPK subunit altered circadian locomotor rhythms in *Drosophila*

AMPK functions as a holoenzyme consisting of a catalytic subunit (α subunit) and two regulatory subunits (β and γ subunits). In mammals, the α subunit is encoded by two isoforms, the β

subunit is encoded by two isoforms, and the γ subunit is encoded by three isoforms. In contrast, in *Drosophila*, each subunit is encoded by one isoform; *snf1A* encodes the α subunit, *alicorn* encodes the β subunit, and *snf4A γ* encodes the γ subunit (Pan and Hardie, 2002). To examine the *in vivo* relevance of the interaction between the AMPK γ subunit and CLK, we analyzed the locomotor behavior of flies in which expression of each AMPK subunit was downregulated in *timeless*-expressing cells using a binary GAL4/UAS system (Brand and Perrimon, 1993). To enhance the RNA interference efficiencies, we coexpressed the UAS-*dicer2* (UAS-*d2*) transgene. The circadian locomotor rhythms of knockdown flies were compared with two lines of control flies, *tim-Gal4; d2* crossed to *w¹¹¹⁸* (*tim>d2*) and UAS-AMPK α ^{Ri} (Table 1). During the LD cycle, control flies exhibited a typical bimodal locomotor activity at times that demonstrated anticipation of the light-on and light-off periods (Fig. 2A, top), which subsequently persisted under DD (Fig. 2A, bottom). Control flies exhibited 23.7–24.5 h period with 30%–98% rhythmicity depending on the lines. In AMPK α knockdown (*tim>d2*, AMPK α ^{Ri-1} and *tim>d2*, AMPK α ^{Ri-2}, named AMPK α Ri-1 and AMPK α Ri-2, respectively) and AMPK β knockdown (*tim>d2*, AMPK β ^{Ri-1} and *tim>d2*, AMPK β ^{Ri-2}, named AMPK β Ri-1 and AMPK β Ri-2, respectively) flies, the morning peak was similar to the peak in the control flies during the LD cycle, but the phase of the evening locomotor peak was slightly delayed (Fig. 2B, top). Consistent with a long periodicity, both morning and evening locomotor activities were delayed under DD (Fig. 2B, bottom). The phase of evening locomotor activities was slightly advanced in AMPK γ knockdown (*tim>d2*, AMPK γ ^{Ri-1} and *tim>d2*, AMPK γ ^{Ri-2}, named AMPK γ Ri-1 and AMPK γ Ri-2, respectively) flies under the LD cycle (Fig. 2B, top), and bimodal locomotor activity was absent under DD (Fig. 2B, bottom). AMPK α Ri-1 and AMPK α Ri-2 flies exhibited a $\sim$ 1.5-h-long period with significantly reduced rhythmicity compared with control flies. AMPK β Ri-1 flies exhibited a 1-h-long period with strong rhythmicity, and the AMPK β Ri-2 flies exhibited a 2.5-h-long period with slightly reduced rhythmicity compared with control flies. AMPK γ Ri-1 and AMPK γ Ri-2 flies were arrhythmic, indicating that knockdown of AMPK γ led to the most pronounced phenotype among the AMPK subunit knockdown flies.

Table 1. Behavior analysis of AMPK subunit knockdown flies^a

Genotype	Temperature (°C)	Number ^b	Rhythmicity ^c (%)	Tau	Power ^d
<i>tim>dicer2</i>	25	32	93.8	24.3 ± 0.07	106.7 ± 7.84
UAS-AMPK α^{Ri} –1 ^e	25	47	91.5	23.7 ± 0.05	144 ± 9.37
<i>tim>dicer2</i> , AMPK α^{Ri} –1	25	31	45.2	25.9 ± 0.2 ^{k,l}	42.2 ± 5.45
UAS-AMPK α^{Ri} –2 ^f	25	45	55.6	24.1 ± 0.05	113.9 ± 10.69
<i>tim>dicer2</i> , AMPK α^{Ri} –2	25	32	9.7	25.5 ± 0.23 ^{k,l}	75.3 ± 14.27
UAS-AMPK β^{Ri} –1 ^g	25	48	75	24.2 ± 0.07	147.6 ± 12.96
<i>tim>dicer2</i> , AMPK β^{Ri} –1	25	32	90.6	25.3 ± 0.45	161 ± 14.38
UAS-AMPK β^{Ri} –2 ^h	25	47	97.9	23.7 ± 0.03	274.5 ± 11.51
<i>tim>dicer2</i> , AMPK β^{Ri} –2	25	32	84.4	26.8 ± 0.15 ^{k,l}	180.5 ± 11.96
UAS-AMPK γ^{Ri} –1 ⁱ	25	39	89.7	24.5 ± 0.07	132 ± 10.26
<i>tim>dicer2</i> , AMPK γ^{Ri} –1	25	31	3.2	Arrhythmic	NA
UAS-AMPK γ^{Ri} –2 ^j	25	41	29.3	24.2 ± 0.07	90.4 ± 14.5
<i>tim>dicer2</i> , AMPK γ^{Ri} –2	25	29	0	Arrhythmic	NA
<i>tub-G80^{TS}</i> , <i>tim>dicer2</i>	18	31	80.6	23.8 ± 0.1	94.6 ± 7.09
<i>tub-G80^{TS}</i> , <i>tim>dicer2</i>	29	30	96.7	23.2 ± 0.41	206.2 ± 13.32
<i>tub-G80^{TS}</i> , <i>tim>dicer2</i> , AMPK α^{Ri} –2	18	30	56.7	24.1 ± 0.15	86.2 ± 7.45
<i>tub-G80^{TS}</i> , <i>tim>dicer2</i> , AMPK α^{Ri} –2	29	29	86.2	24.5 ± 0.06 ^m	180.2 ± 15.34
<i>tub-G80^{TS}</i> , <i>tim>dicer2</i> , AMPK β^{Ri} –2	18	31	54.8	24.2 ± 0.26	94.9 ± 8.85
<i>tub-G80^{TS}</i> , <i>tim>dicer2</i> , AMPK β^{Ri} –2	29	15	100	25.1 ± 0.11 ^m	188.7 ± 13.7
<i>tub-G80^{TS}</i> , <i>tim>dicer2</i> , AMPK γ^{Ri} –2	18	31	45.2	24.3 ± 0.09	76.2 ± 6.63
<i>tub-G80^{TS}</i> , <i>tim>dicer2</i> , AMPK γ^{Ri} –2	29	13	15.4	23.5 ± 0.25 ^m	50.2 ± 4.55
<i>pdf>dicer2</i>	25	30	93.3	24.5 ± 0.09	156 ± 12.84
<i>pdf>dicer2</i> , AMPK β^{Ri} –2	25	31	93.5	26.8 ± 0.18 ⁿ	127.7 ± 8.7
<i>pdf>dicer2</i> , <i>Clk</i>	25	30	73.3	23.5 ± 0.22 ⁿ	109 ± 10.65
<i>pdf>dicer2</i> , AMPK β^{Ri} –2, <i>Clk</i>	25	14	64.3	24 ± 0.74 ⁿ	77.3 ± 11.47

^aFlies were maintained at the indicated temperatures and exposed to 4 d of 12 h:12 h LD followed by 6–8 d of DD.^bTotal number of flies that survived until the end of the testing period.^cPercentage of flies with activity rhythms exhibiting a power value ≥ 10 and a width value ≥ 2 .^dRelative measure of the robustness of the rhythm.^eVDR1827.^fNIG3051R-1.^gNIG8057R-2.^hNIG8057R-4.ⁱNIG17299R-3.^jNIG17299R-1.^kSignificantly different from the UAS-Ri control under the same conditions (one-way ANOVA, Tukey's test, $p < 0.05$).^lSignificantly different from the *tim>dicer2* control under the same conditions (one-way ANOVA, Tukey's test, $p < 0.05$).^mSignificantly different from the 18°C case in the same genetic background (Student's *t* test, $p < 0.05$).ⁿTwo-way ANOVA showed significant differences in the AMPK β knockdown ($F_{(1,84)} = 26.23, p = 1.83 \times 10^{-6}$), *Clk* overexpression ($F_{(1,84)} = 120.03, p = 5.40 \times 10^{-18}$), and interaction ($F_{(1,84)} = 37.20, p = 2.95 \times 10^{-8}$).

The total daily locomotor activity counts of most knockdown flies did not exhibit significant reduction compared with parental UAS-AMPK α^{Ri} control flies, although those of AMPK α Ri-1 and AMPK β Ri-1 flies were reduced (Fig. 2C). Because the reduction was observed in some, but not all, lines of knockdown flies and the extent of reduction was not severe, we reasoned that downregulation of AMPK only in *timeless*-expressing cells did not lead to gross locomotor defects, whereas a previous report showed that pan-neuronal AMPK γ inactivation leads to severe locomotor defects (Nagarkar-Jaiswal et al., 2017).

Reduced expression of AMPK α , AMPK β , and AMPK γ mRNA was verified in AMPK α Ri-2, AMPK β Ri-2, and AMPK γ Ri-2 flies, respectively, by real-time qRT-PCR (Fig. 2D–F). The mRNA levels of the AMPK α , β , and γ subunits did not exhibit diurnal oscillations in control flies and were significantly reduced in each knockdown fly at both ZT8 and ZT20. The AMPK α protein level was not reduced in whole-head extracts of AMPK α Ri flies, probably because gene knockdown in AMPK α Ri flies was limited to *tim*-expressing cells (data not shown). Nonetheless, reduction of AMPK α protein level was confirmed in AMPK α knockdown larvae, in which expression of AMPK α^{Ri} –1 or AMPK α^{Ri} –2 was driven by actin-Gal4 (Fig. 2G,H).

AMPK subunit downregulation affected sLN ν integrity and *per* expression in clock neurons

To understand the circadian rhythm defects in AMPK subunit knockdown flies, we inspected PDF neurons, which play major roles in maintaining locomotor rhythms (Helfrich-Förster, 1998; Renn et al., 1999; Nitabach et al., 2002). In AMPK γ Ri-2 flies, which showed the most severe rhythm defects compared with AMPK α or AMPK β Ri-2 flies, PDF immunostaining at ZT2 revealed that the neurites that project dorsally from the sLN ν to DN ν s were missing, whereas these neurites were present and produced a strong signal in control flies (Fig. 3A). The morphology of the neurites that projected dorsally from the sLN ν was somewhat defective in AMPK α Ri-2 flies, but not in AMPK β Ri-2 flies (Fig. 3A). To investigate whether the absence of PDF-stained dorsally projecting neurites was a consequence of low PDF expression or loss of sLN ν neurons, we observed LN ν neurons directly using the reporter protein nls-GFP, which allows visualization of the soma, and mCD8::GFP, which allows visualization of neurites under the control of the *pdf*-Gal4 driver. Although strong GFP signals were observed in the ILN ν s of control and AMPK γ Ri-2 flies, the GFP signals in the sLN ν s of AMPK γ Ri-2 flies were very weak or absent compared with the signals in

Figure 2. Circadian locomotor behaviors were altered in AMPK subunit knockdown flies. **A, B**, Daily activity profiles of young adult male flies on the last day of LD (top) representing a given genotype: tim-Gal4>UAS-dicer2 (tim>d2), tim-Gal4>UAS-dicer2, UAS-AMPK α^{RI-1} (tim>d2, AMPK α^{RI-1}), tim-Gal4>UAS-dicer2, UAS-AMPK α^{RI-2} (tim>d2, AMPK α^{RI-2}), tim-Gal4>UAS-dicer2, UAS-AMPK β^{RI-1} (tim>d2, AMPK β^{RI-1}), tim-Gal4>UAS-dicer2, UAS-AMPK β^{RI-2} (tim>d2, AMPK β^{RI-2}), tim-Gal4>UAS-dicer2, UAS-AMPK γ^{RI-1} (tim>d2, AMPK γ^{RI-1}), tim-Gal4>UAS-dicer2, UAS-AMPK γ^{RI-2} (tim>d2, AMPK γ^{RI-2}) are shown. Bottom, Actograms of flies during 5–7 consecutive days of DD. To better visualize the rhythmic behavior, each row in an actogram was double-plotted. Light gray bars represent day-time activity. Dark gray bars represent subjective day-time activity. Black bars represent night-time and subjective night-time activity. Hatched horizontal bar represents subjective day. Black horizontal bar represents subjective night. ZT, Zeitgeber time. ZT0 is light-on time. CT, Circadian time. CT0 is subjective day start time. **C**, Total number of beam crossings of the single fly during the last day of LD was calculated and averaged for each given genotype of flies. Values indicate mean $\pm$ SEM. Statistically significant differences in the total activity counts between UAS-AMPK α^{RI} control flies and knockdown flies (Student's *t* test): **p* < 0.05; ****p* < 0.01. **D–F**, Flies from the following strains were collected at ZT8 and ZT20: tim>d2; tim>d2, AMPK α^{RI-2} ; tim>d2, AMPK β^{RI-2} ; and tim>d2, AMPK γ^{RI-2} . Total RNA was extracted, and mRNA levels of AMPK α (**D**), AMPK β (**E**), and AMPK γ (**F**) were quantified using real-time qRT-PCR. The mRNA levels in tim>d2 at ZT8 were set to 1, and the other values were normalized to these values. Values indicate mean $\pm$ SEM of three independent (Figure legend continues.)

Figure 3. AMPK γ Ri flies lack sLN_vs and dorsal projections from sLN_vs. **A**, Brains from the following fly strains were dissected at ZT2 and stained with anti-PDF (C7): *tim>d2*; *tim>d2, AMPKα^{Ri-2}*; *tim>d2, AMPKβ^{Ri-2}*; *tim>d2, AMPKγ^{Ri-2}*; *pdf>nls-GFP*; *pdf>nls-GFP, AMPKγ^{Ri-2}*; *pdf>mCD8::GFP*; *pdf>mCD8::GFP, AMPKγ^{Ri-2}*. Cell bodies (**B**) or neurites (**B**) of LN_vs were visualized by nls-GFP or mCD8::GFP expression under the control of pdf-Gal4. **C**, POT, Posterior optic tract. Arrowheads indicate dorsally projecting neurites from sLN_v.

the control flies (Fig. 3B). In addition, dorsally projecting neurites that expressed mCD8::GFP were clearly visible in control flies but were absent in AMPK γ Ri-2 flies (Fig. 3C). Thus, knock-down of AMPK γ severely impaired the neuronal integrity of sLN_v neurons but not ILN_v neurons. Previous reports have shown that the anticipation of light-on and light-off during LD is primarily controlled by ILN_v neuronal activity (Cusumano et al., 2009; Sheeba et al., 2010). Further, it has been shown that sLN_v neurons are required to synchronize the clock neuronal network to maintain the circadian rhythmicity under DD (Renn et al., 1999; Peng et al., 2003; Lin et al., 2004). In AMPK γ Ri flies, quasi-normal daily locomotor activity during LD and complete loss of rhythmicity under DD could be attributed to intact ILN_v neurons and impaired sLN_v neurons.

The morphological defects observed in the sLN_vs of AMPK γ Ri-2 flies led us to examine whether the circadian behavior phenotypes manifested by AMPK downregulation are caused by developmental effects. To test this, we used the TARGET system with tubulin-Gal80^{ts} (McGuire et al., 2004). Flies were raised at the permissive temperature (18°C) at which Gal80^{ts} inhibited expression of UAS-AMPK^{Ri}. Then, locomotor behavior analysis was conducted at the permissive temperature and the restrictive temperature (29°C). At 29°C, Gal80^{ts} does not function and Gal4 activates the expression of UAS-AMPK^{Ri}. In control flies, the

period was shortened by ~ 0.6 h with enhanced rhythmicity at 29°C compared with the period and rhythmicity at 18°C, although the period difference was not statistically significant (Table 1). The periods in AMPK α ^{Ri-2}, AMPK β ^{Ri-2}, and AMPK γ ^{Ri-2} flies were ~ 24 h at 18°C. At 29°C, the periods in AMPK α ^{Ri-2} and AMPK β ^{Ri-2} flies were lengthened by ~ 0.5 h and ~ 1 h, respectively, and rhythmicity was enhanced, whereas the period in AMPK γ ^{Ri-2} flies was shortened by ~ 0.8 h and rhythmicity was greatly reduced. The period alterations between 18°C and 29°C in AMPK α ^{Ri-2}, AMPK β ^{Ri-2}, and AMPK γ ^{Ri-2} flies were statistically significant. Thus, behavioral analysis demonstrates that the depletion of AMPK α and AMPK β subunits in adult flies in an acute manner appears to be sufficient to lengthen the period, and depletion of the AMPK γ subunit sufficient to reduce rhythmicity. Compared with life-long knockdowns, we observed weaker effects on period and rhythmicity, probably because the duration of AMPK^{Ri} expression was shorter; and thereby, the knockdown efficiency was lower. These data indicate that the circadian rhythm phenotypes manifested by downregulation of AMPK subunits are due to active requirements in clock cells and not to secondary effects on development.

Next, we performed PER immunostaining at ZT2 in control and AMPK subunit knockdown flies (Fig. 4A–D). PER-stained sLN_vs were missing in AMPK α Ri-2 and AMPK γ Ri-2 flies, and the dorsally projecting neurites from the sLN_vs were absent in AMPK γ Ri-2 flies and were fragmented in AMPK α Ri-2 flies. On the other hand, sLN_vs and the dorsally projecting neurites from the sLN_vs were intact in the AMPK β Ri-2 flies. PER levels were quantified in ILN_vs, LN_ds, and DN₁s (Fig. 4E–G). In all AMPK subunit knockdown flies, PER levels were reduced in ILN_vs, LN_ds, and DN₁s compared with control flies, although some differences were not statistically significant. It appears that the morphological defects in the sLN_vs were most severe in the γ knockdown,

(Figure legend continued.) experiments. Student's *t* test: **p* < 0.05; ***p* < 0.01; ****p* < 0.001; *****p* < 0.0001. **G**, Protein extracts were prepared from larvae of the indicated genotypes at ZT8 and ZT20 and were analyzed by immunoblotting with anti-AMPK α antibody. β -Tubulin served as the loading control. A representative image from three independent experiments is shown. **H**, Quantification of AMPK α protein levels was performed with ImageJ. The AMPK α level in *actin>d2* at ZT8 was set to 1, and the other values were normalized to this value. Values indicate mean $\pm$ SEM of three independent experiments. One-way ANOVA with Tukey's test: **p* < 0.05; ***p* < 0.01; ****p* < 0.001.

Figure 4. PER levels were reduced in pacemaker neurons of AMPK subunit knockdown flies. **A, B, C, D**, Brains from flies of the indicated genotypes were dissected at ZT2 and stained with anti-PER (Rb1, green) and anti-PDF (C7, red) antibodies. The dorsally projecting neurites from sLN_vs (arrowheads) showed morphological defects (**B**) or were missing (**D**). **E, F, G**, PER fluorescence intensities in ILN_v, LN_ds, and DN₁ neurons were quantified. Values indicate mean $\pm$ SEM ($n = 11$ –35). One-way ANOVA with Tukey's test: * $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$; **** $p < 0.0001$.

Figure 5. Transcription of CLK-target core clock gene is reduced in AMPK β Ri flies. **A, B**, Flies of the indicated genotype were collected during LD and total RNA was isolated. Pre-mRNA levels of *per* (**A**) and *vri* (**B**) were quantified using real-time qRT-PCR. The pre-mRNA levels in *tim>d2* at ZT4 were set to 1, and the other values were normalized to these values. Values indicate mean $\pm$ SEM of three independent experiments. * $p < 0.05$ (Student's *t* test). **C, D**, Flies of the indicated genotypes were collected at ZT2 (**C**) and ZT17 (**D**) during LD. The brains were stained with anti-PER (Rb1, green), anti-VRI (gp2, green), and anti-PDF (red) antibodies. Representative images of sLN_vs and LN_ds from two independent experiments are shown. PER and VRI fluorescence intensities were quantified. Values indicate mean $\pm$ SEM ($n = 16$ –56). * $p < 0.05$ (Student's *t* test). *** $p < 0.001$ (Student's *t* test).

then the α knockdown, and were the least severe in the β knockdown; thus, the highly arrhythmic locomotor behaviors in the α and γ Ri flies were likely caused by damaged sLN_vs. Nonetheless, reduced PER levels in clock neurons were observed in all three AMPK subunit knockdown flies. Our data suggest that AMPK enzymatic activity is required to maintain the integrity and cellular oscillator function of sLN_v neurons. Because sLN_vs are required for synchronizing circadian neural network for locomotor rhythm, we decided to use the AMPK β knockdown flies containing intact sLN_vs to further examine the biochemical function of AMPK in the cellular oscillator.

AMPK β knockdown reduced CLK-driven transcription by reducing CLK levels

Intriguingly, sLN_vs and dorsal projecting neurites are missing in *Clk* and *cyc* mutants (Blau and Young, 1999; Park et al., 2000; Allada et al., 2003), phenocopying the situation when AMPK α or AMPK γ was downregulated (Fig. 3). In addition, because AMPK interacted with CLK (Fig. 1), we reasoned that CLK/CYC activity may be affected in AMPK subunit knockdown flies. To examine whether CLK/CYC-mediated transcription was affected in AMPK β subunit knockdown flies, we determined the pre-mRNA levels of *per* and *vri*, which are core clock genes that are targeted by CLK/CYC (Fig. 5A,B). It has been established that pre-mRNA levels can be used as a read-out of transcription (Menet et al., 2010; Luo et al., 2012). Both *per* and *vri* pre-mRNA levels were reduced in AMPK β subunit knockdown flies, indicating that downregulation of AMPK β reduced CLK/CYC-dependent transcription. Next, we performed immunostaining to examine PER and VRI protein levels at their peak time points: that is, ZT2, and ZT17, respectively, in sLN_vs and LN_ds (Fig. 5C,D). Consistent with the reduction in CLK/CYC-dependent transcription, we found that PER and VRI protein levels were reduced. The re-

duced PER and VRI protein levels may have impeded the pace of the circadian clock in AMPK β Ri-2 flies.

To determine whether AMPK directly phosphorylates CLK, we performed an *in vitro* kinase assay using purified AMPK holoenzyme and bacterially expressed GST-tagged CLK as the substrate. We found that AMPK phosphorylates CLK and that the phosphorylation was greatly enhanced by the addition of AMP (Fig. 6A). To assess how AMPK-mediated phosphorylation affects CLK, we examined daily CLK profiles. Hyperphosphorylated isoforms of CLK were observed in the early-day and late-night periods (ZT4, ZT20, and ZT23.8), and hypophosphorylated isoforms of CLK were observed during mid-day (ZT8 to ZT16) (Fig. 6B). A side-by-side comparison of CLK showed no significant differences in phosphorylation pattern in control and AMPK β Ri-2 flies. In addition, CLK protein levels in whole-head extracts from control and AMPK β knockdown flies were similar throughout the day (Fig. 6C). Nevertheless, we reasoned that the effect of AMPK on CLK protein may be different in clock cells, and we tested this by immunostaining. Indeed, CLK levels were lower in sLN_vs and LN_ds in AMPK β knockdown flies than in control flies (Fig. 6D). Moreover, both PER and CLK levels in sLN_vs were lower in flies in which AMPK β was downregulated in an acute manner via the Gal80^{TS} TARGET system than in control flies (Fig. 6E). Collectively, these observations indicate that AMPK-mediated phosphorylation stabilizes CLK in pacemaker neurons and enhances expression of CLK/CYC target genes.

Expression of CLK rescued the arrhythmicity caused by the AMPK β knockdown

If AMPK downregulation slowed the clock by affecting CLK levels, overexpression of CLK may rescue the circadian locomotor rhythm defect caused by AMPK β knockdown. Because overexpression of *Clk* in combination with the *tim*-GAL4 driver is de-

Figure 6. AMPK directly phosphorylated CLK *in vitro* and stabilized CLK in clock neurons. **A**, GST-CLK and GST purified from *E. coli* were incubated with AMPK holoenzyme and γ - 32 P-ATP with increasing amounts of AMP. The samples were then subjected to SDS-PAGE, and the proteins were transferred to PVDF membranes. Following Ponceau S staining, phosphorylation of GST-CLK and GST was visualized by autoradiography. **B**, Protein extracts from the heads of control (Ct) and AMPK β knockdown flies (Ri) were prepared at the indicated ZT and were analyzed by immunoblotting with anti-CLK (gp208) antibody. Actin served as the loading control. **C**, Relative levels of CLK were determined by measuring band intensities using ImageJ software. Values indicate mean $\pm$ SEM ($n = 3$). **D**, Flies of the indicated genotypes were collected at ZT24 during LD, and the brains were stained with antibodies to CLK (gp50, green) and PDF (C7, red). Representative images of sLN_vs and LN_vs from two independent experiments are shown. CLK fluorescence intensities were quantified. Values indicate mean $\pm$ SEM ($n = 24$ –53). Statistically significant differences between control and AMPK β Ri flies (Student's *t* test): ** $p < 0.01$, *** $p < 0.001$, **** $p < 0.0001$. **E**, Flies of the indicated genotypes were collected at ZT2 at 29°C. The brains were stained with antibodies to CLK (gp50, green) and PDF (C7, red) or PER (Rb1, green) and PDF (C7, red). Representative images of sLN_vs are shown. CLK and PER fluorescence intensities were quantified. Values indicate mean $\pm$ SEM ($n = 24$ –37). Statistically significant differences between control and AMPK β Ri flies (Student's *t* test): ** $p < 0.01$, *** $p < 0.001$.

developmentally lethal (Zhao et al., 2003), we expressed UAS-*Clk* under control of the pdf-GAL4 driver. We confirmed overexpression of CLK by immunostaining sLN_vs (Fig. 7A,B). Knockdown of AMPK β by the pdf-GAL4 driver in LN_vs was sufficient to cause long periods similar to the AMPK β knockdown in the *tim*-expressing cells and to downregulate CLK levels in sLN_vs (Table 1; Fig. 7D). Overexpression of CLK suppressed the long-period phenotype of the AMPK β knockdown and restored the 24 h period (Fig. 7C). These results indicate that the key mechanism underlying the circadian locomotor defect caused by AMPK β knockdown in flies is downregulation of CLK levels.

Discussion

Daily rhythmic environmental changes entrain the circadian clock system of an organism to manifest the appropriate physiology and behavior on a circadian time-scale. In eukaryotes, oscillation of gene expression driven by circadian transcription factors is the basic framework for the cell-autonomous oscillator (for reviews, see Dunlap and Loros, 2004; Hardin, 2005; Takahashi, 2017). Thus, regulation of circadian transcription factors is fundamental to synchronizing the clock system with environmental changes. Here, we report that the AMPK γ subunit copu-

rified with CLK in *Drosophila* S2 cells and that AMPK activity is required for circadian locomotor rhythms because it enhances CLK-dependent transcriptional activation in the core clock neurons of *Drosophila*.

AMPK functions as a holoenzyme consisting of a catalytic α subunit and two regulatory subunits, β and γ . In mammals, full activation of AMPK occurs upon phosphorylation of Thr172 (analogous to Thr184 in *Drosophila*) of the α subunit by upstream kinases, such as LKB and CAMKK2, which are activated by energy stress and an increase in the intracellular Ca^{2+} level, respectively (for review, see Ross et al., 2016; Garcia and Shaw, 2017). AMP binding to the γ subunit prolongs Thr172 phosphorylation of the α subunit, thereby enhancing AMPK activity (Xiao et al., 2011; Gowans et al., 2013). The β subunit plays a stoichiometric and stabilizing role in the trimeric complex by binding the α and γ subunits in a 1:1:1 ratio (Woods et al., 1996). Binding of a novel AMPK-activating compound to the AMPK β subunit protects the AMPK α subunit Thr172 from dephosphorylation, indicating that the AMPK β subunit also regulates the phosphorylation status of AMPK α (Göransson et al., 2007; Sanders et al., 2007). Consistently, mutant mice lacking AMPK β 1 exhibit severe brain abnormalities caused by reduced AMPK activity (Dasgupta and Milbrandt, 2009). Consequently, in

Figure 7. Expression of *Clk* rescued the long period and arrhythmicity induced by AMPK β knockdown. **A, B**, Flies of the indicated genotypes were collected at ZT2 during LD. The brains were stained with anti-CLK (gp50, green) and anti-PDF (red) antibodies. Representative images of sLN_vs are shown. CLK fluorescence intensities were quantified. Values indicate mean $\pm$ SEM ($n = 18–21$). One-way ANOVA with Dunnett's test: $*p < 0.05$; $****p < 0.0001$. **C**, Graph represents the period lengths in *pdf>d2* and *pdf>d2*, AMPK β ^{RI} flies without (–) or with (+) *Clk* overexpression by *pdf*-Gal4 driver. Values indicate mean $\pm$ SEM. Two-way ANOVA showed significant differences in the AMPK β knockdown ($F_{(1,84)} = 26.23$, $p = 1.83 \times 10^{-6}$), *Clk* overexpression ($F_{(1,84)} = 120.03$, $p = 5.40 \times 10^{-18}$), and interaction ($F_{(1,84)} = 37.20$, $p = 2.95 \times 10^{-8}$). **D**, Daily activity profiles of young adult male flies of the indicated genotypes on the last day of LD (top) and on the first day of DD (middle) are shown. Bottom, Actograms of flies during 7 consecutive days of DD. To better visualize the rhythmic behavior, each row in an actogram was double-plotted. Light gray bars represent day-time activity. Dark gray bars represent subjective day-time activity. Black bars represent night-time and subjective night-time activity. Hatched horizontal bar represents subjective day. Black horizontal bar represents subjective night. ZT, Zeitgeber time. ZT0 is light-on time. CT, Circadian time. CT0 is subjective day start time.

this study, we reasoned that downregulation of expression of each AMPK subunit would reduce the activity of the AMPK holoenzyme. Our results showed that knockdown of each AMPK subunit produced circadian locomotor rhythm defects, supporting the notion that AMPK activity is required for the circadian rhythm in *Drosophila*.

In mammals, AMPK affects the core clock by regulating circadian repressor proteins. AMPK phosphorylates mCRY1 and mCRY2, which stimulates their interaction with F-box/LRR-repeat protein 3 (FBXL3), an E3 ligase, leading to degradation of CRY (Lamia et al., 2009). AMPK has also been shown to phosphorylate serine 389 of CK1 ϵ , which enhances CK1 ϵ enzymatic activity and degradation of mPER2 (Um et al., 2011). Collectively, in mammals, AMPK destabilizes circadian repressor proteins and induces derepression of CLK/BMAL1-dependent transcription. The role of AMPK in regulating expression of clock genes was reported in skeletal muscle cells following treatment with 5-aminoimidazole-4-carboxamide ribonucleotide, a stimulator of AMPK activity, albeit the detailed molecular mechanism was not provided (Vieira et al., 2008).

In this study, we propose that, in *Drosophila*, AMPK regulates the core clock by increasing the activity and level of CLK. This

proposition is based on several observations. First, if AMPK phosphorylates and destabilizes CRY as in mammals, then downregulation of AMPK subunits would result in increased CRY levels in *Drosophila*. However, previous studies have shown that overexpression of *cry* under the control of a *tim*-GAL4 or a *pdf*-GAL4 driver did not affect behavioral rhythms in *Drosophila* (Dissel et al., 2004; Collins et al., 2006). Thus, AMPK does not appear to affect the core clock by regulating CRY stability in *Drosophila*. Second, *per* and *vri* pre-mRNA levels were lower in AMPK β knockdown flies than in control flies (Fig. 5), indicating that CLK/CYC-dependent transcription requires AMPK activity (Menet et al., 2010; Luo et al., 2012; Andreatza et al., 2015). We also showed that PER and VRI protein levels were lower in pacemaker neurons of AMPK β knockdown flies than in the control flies, which is consistent with our pre-mRNA results. Furthermore, rescue of locomotor behavior in AMPK β knockdown flies by LN_v-specific CLK expression strongly supports the notion that compromised CLK/CYC transcription underlies the circadian rhythm defect that occurs when AMPK is downregulated. Third, we showed that purified CLK protein is directly phosphorylated by AMPK in an AMP-dependent manner using the *in vitro* kinase assay (Fig. 6A). To identify the AMPK phosphorylation site(s) on

CLK, we searched candidate phosphosites using Scansite 4.0 (<https://scansite4.mit.edu/4.0/#scanProtein>). We mutated 10 Ser/Thr residues in CLK to alanine and used the *in vitro* kinase assay to evaluate phosphorylation of these CLK mutants. We did not observe any significant differences in the phosphorylation of these mutants compared with WT CLK (data not shown), perhaps because the predicted phosphosites were not the actual phosphosites or because alteration of a single residue is not enough to affect the degree of phosphorylation *in vitro*. Further studies are needed to determine the AMPK phosphorylation site(s) in CLK. Finally, the impaired integrity of the sLN_vs and the dorsal projections observed in AMPK α and AMPK γ *Ri* flies matched the phenotype of flies with reduced CLK and CYC activities (Fig. 3; and see below). Thus, we conclude that AMPK regulates the core clock in *Drosophila* by activating CLK, a positive component of the clock. It is intriguing to note that, while the target of AMPK action is different in *Drosophila* and mammals (positive element vs negative element, respectively), the outcome is in the same direction (i.e., the enhancement of circadian transcription). Nonetheless, there is also possibility that other circadian clock protein(s) are affected by AMPK and regulate circadian rhythm.

Downregulation of the AMPK α or the AMPK γ subunit in *tim*-expressing cells led to arrhythmicity accompanied with the disruption of sLN_v cell bodies and dorsally projecting neurites. It has been shown that sLN_vs and dorsally projecting neurites are absent when CLK and CYC activities are compromised in *Clk*^{Jrk}, *Clk*^{ar}, *Clk*^{out}, and *cyc*⁰ mutants (Blau and Young, 1999; Park et al., 2000; Allada et al., 2003). In addition, it has recently been shown that overexpression of the dominant negative form of *microtubule star*, which encodes a catalytic subunit of PP2A, in *tim*-expressing cells induces morphological defects in sLN_vs (Andreazza et al., 2015). Although the authors of the study did not discuss the possible underlying mechanism, because PP2A controls the phosphorylation status of CLK, which is critical for CLK stability and activity, dysregulation of CLK may underlie the defects in the sLN_vs in the *microtubule star* mutant (Kim and Ederly, 2006; Andreazza et al., 2015). Furthermore, VRI, the downstream target of CLK/CYC, controls the accumulation of the neuropeptide PDF in a post-transcriptional manner (Blau and Young, 1999; Gunawardhana and Hardin, 2017). Thus, dysregulation of CLK/CYC activity may underlie the impaired integrity of sLN_vs in AMPK α and AMPK γ *Ri* flies, which is consistent with the observation that CLK is required for sLN_v development and integrity (Lerner et al., 2015). Interestingly, transcriptome analysis revealed that *Clk* expression is reduced in human epileptogenic tissue, and a reduction in *Clk* expression contributes to malformation of the dendritic spine. These data suggest that CLK transcriptional activity is required for neuronal structural integrity (Li et al., 2017).

ATP levels and Ca²⁺ levels, which are upstream signals of AMPK, have been shown to display daily rhythms in several tissues and organisms (Yamazaki et al., 1994; Ikeda et al., 2003; Enoki et al., 2012; Fustin et al., 2012; Goya et al., 2016). AMPK activity has also been shown to exhibit a daily rhythm in the hypothalamus, retina, liver, and fibroblasts in a cell-autonomous manner (Um et al., 2011; Barnea et al., 2012; Huang et al., 2015). The differences in the active phases and regulatory mechanisms in the tissues and cells suggest that daily AMPK rhythms may be controlled in a tissue- and cell-specific manner. We also sought to explore whether AMPK activity might show daily rhythm, coupled with regulation of CLK levels in *Drosophila*. Nonetheless, the

mRNA levels of the AMPK α , β , and γ subunits did not show diurnal oscillations (Fig. 2D–F, compare ZT8 and ZT20). Because the available anti-AMPK α antibody does not work in the immunostaining assay, Western blot analysis of whole fly head extracts was performed; no rhythms in daily AMPK α protein levels were observed (data not shown). However, the possibility that AMPK α levels oscillate in a cell-specific manner cannot be ruled out. Regarding AMPK activity, a widely used AMPK α Thr172 phosphospecific antibody could not reliably determine the phosphorylation status of *Drosophila* AMPK α Thr184 (data not shown). Thus, currently, it is not elucidated whether the daily rhythmic AMPK activity and/or levels are coupled to regulate rhythmic CLK/CYC-dependent transcription in a daily basis in *Drosophila*. Nevertheless, pronounced acute metabolic stress or an increase in the intracellular Ca²⁺ level can induce AMPK activation and enhance CLK/CYC-dependent transcription to reset the phase.

In *Drosophila*, the crosstalk mechanism between nutrients, energy metabolism, and the core clock has been elucidated. The nutrient signal regulator AKT and the TOR-S6K pathway determine the circadian period by regulating shaggy/glycogen synthase kinase 3 β , thereby regulating nuclear entry of TIM (Zheng and Sehgal, 2010). O-GlcNAc modification of PER, which may be under the control of cellular glucose metabolism, sets the circadian period by regulating nuclear entry of PER (Kim et al., 2012; Kaasik et al., 2013). In this study, we showed that AMPK, which is sensitive to the AMP/ATP ratio and the internal Ca²⁺ concentration, regulates the circadian rhythm by regulating the level and activity of CLK.

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