

Circadian Clocks and Feeding Time Regulate the Oscillations and Levels of Hepatic Triglycerides

Yaarit Adamovich,¹ Liat Rousso-Noori,¹ Ziv Zwighaft,¹ Adi Neufeld-Cohen,¹ Marina Golik,¹ Judith Kraut-Cohen,¹ Miao Wang,² Xianlin Han,² and Gad Asher^{1,*}

¹Department of Biological Chemistry, Weizmann Institute of Science, Rehovot 76100, Israel

²Diabetes and Obesity Research Center, Sanford-Burnham Medical Research Institute, Orlando, FL 32827, USA

*Correspondence: gad.asher@weizmann.ac.il

<http://dx.doi.org/10.1016/j.cmet.2013.12.016>

SUMMARY

Circadian clocks play a major role in orchestrating daily physiology, and their disruption can evoke metabolic diseases such as fatty liver and obesity. To study the role of circadian clocks in lipid homeostasis, we performed an extensive lipidomic analysis of liver tissues from wild-type and clock-disrupted mice either fed ad libitum or night fed. To our surprise, a similar fraction of lipids (~17%) oscillated in both mouse strains, most notably triglycerides, but with completely different phases. Moreover, several master lipid regulators (e.g., PPAR α) and enzymes involved in triglyceride metabolism retained their circadian expression in clock-disrupted mice. Nighttime restricted feeding shifted the phase of triglyceride accumulation and resulted in ~50% decrease in hepatic triglyceride levels in wild-type mice. Our findings suggest that circadian clocks and feeding time dictate the phase and levels of hepatic triglyceride accumulation; however, oscillations in triglycerides can persist in the absence of a functional clock.

INTRODUCTION

Circadian clocks play a principal role in coordinating our daily physiology and metabolism. Animal studies and epidemiological evidence suggest that disturbance of circadian rhythms through environmental and genetic effects can lead to metabolic diseases such as hyperlipidemia, fatty liver, and obesity (Asher and Schibler, 2011; Bass, 2012; Froy, 2010; Green et al., 2008). These observations highlight the central role of circadian regulation in lipid homeostasis. Dyslipidemia and obesity are associated with high morbidity and mortality rates, hence elucidating the mechanisms involved in temporal regulation of lipids is of great interest.

The mammalian circadian clock is comprised of a master pacemaker, located in the brain, that synchronizes subsidiary peripheral oscillators, present in virtually all cells of the body. The master clock is entrained by daily light/dark cycles, whereas feeding time appears to be the dominant timing cue (Zeitgeber) for peripheral clocks. Both the master and

peripheral clocks tick in a self-sustained and cell-autonomous fashion. The currently held molecular model for the generation of circadian rhythms is based on interlocked negative transcription feedback loops that drive daily oscillations of core clock and clock-controlled genes (Brown et al., 2012; Dibner et al., 2010; Feng and Lazar, 2012). Briefly, BMAL1/CLOCK drives the expression of *Period* (i.e., *Per1*, *Per2*, and *Per3*) and *Cryptochrome* (i.e., *Cry1* and *Cry2*) genes. In turn, PER and CRY proteins accumulate and repress the transcription of their own genes. An additional essential feedback loop involves the orphan nuclear receptors of the REV-ERB and ROR families. BMAL1 activates *Rev-erb* transcription, which in turn suppresses *Bmal1* expression (Bugge et al., 2012; Cho et al., 2012; Preitner et al., 2002; Solt et al., 2012).

Extensive transcriptome profiling performed throughout the day in liver and additional peripheral organs has demonstrated the pervasive circadian control of physiology and metabolism (Akhtar et al., 2002; McCarthy et al., 2007; Panda et al., 2002; Storch et al., 2002; Vollmers et al., 2009). These studies have revealed that a substantial fraction (~10%) of all liver mRNAs are expressed in a rhythmic fashion; many of them play a role in metabolic processes, including cholesterol and lipid metabolism. Several enzymes participating in lipid biosynthesis and catabolism are expressed in a daily manner (e.g., cytochrome P450s, HMGCoA reductase, and lipin) (Panda et al., 2002). Additional studies have shown diurnal regulation of triglyceride and cholesterol levels in plasma (Hussain and Pan, 2009). Consequently, various genetic mouse models for disrupted clock exhibit impaired lipid metabolism. *Clock* mutant and *Bmal1* knockout mice develop hyperlipidemia and hepatic steatosis (Shimba et al., 2011; Turek et al., 2005). PER2-deficient mice display altered lipid metabolism (Grimaldi et al., 2010), and ablation of REV-ERBs can lead to hepatic steatosis (Bugge et al., 2012).

Hitherto, circadian research in mammals has largely been focused on dissecting transcription expression profiles of core clock and output genes including genes involved in metabolism and cellular homeostasis. Though these gene expression profiles have hinted that many metabolic pathways and their products are oscillating during the day, direct measurements of metabolites throughout the day are still in their infancy. Ueda and colleagues have quantified the spectra of hundreds of metabolites throughout the day, both in mouse and human plasma samples (Kasukawa et al., 2012; Minami et al., 2009). They successfully established a metabolite

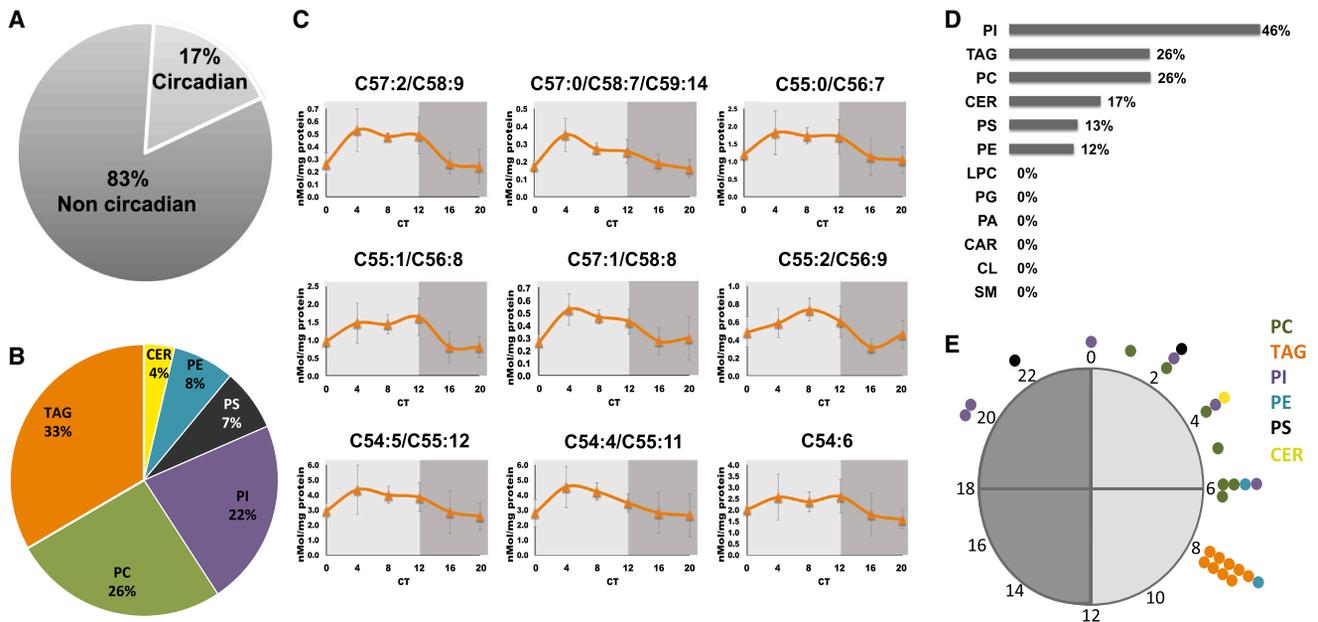


Figure 1. Analysis of Mouse Liver Lipidome

(A) The percentage of lipids found to exhibit a circadian pattern of accumulation in livers of WT mice based on JTK_CYCLE analysis (six time points, $n = 4$ for each, $p < 0.05$). Out of the 159 measured and identified, 27 lipids exhibited circadian pattern of accumulation.

(B) Oscillating lipids species distributed according to their types.

(C) Accumulation profiles of oscillating TAG presented as mean \pm SD.

(D) The percentage of oscillating lipid species within each lipid type.

(E) Daytime distribution of peak phases of oscillating lipids. TAG in orange, PC in green, PI in purple, PS in black, PE in light blue, and CER in yellow. CT0 is when the light was routinely turned on, and CT12 is when the light was turned off in the animal facility. Dark gray represents the subjective night and light gray the subjective day. See also Figure S1.

timetable method to determine internal body time using these profiles. These studies have primarily focused on measuring internal body time by blood metabolomics and to a lesser extent aimed at identifying these metabolites, their metabolic pathways, and circadian clocks dependency. Recent studies performed with human (i.e., blood plasma and saliva) and mouse samples (i.e., liver) identified and measured the levels of about 300 named metabolites throughout the day (Dallmann et al., 2012; Eckel-Mahan et al., 2012). These analyses screened for a variety of primary metabolites (e.g., amino acids, carbohydrates, nucleotides, and lipids). Remarkably, a large fraction of oscillating metabolites identified by these recent reports were lipids.

In this study, we extensively examined the circadian changes in lipid abundance in mouse liver and dissected its clock/feeding dependency. To this aim, we performed a temporal and quantitative lipidomic analysis of livers from wild-type (WT) and clock-disrupted mice (i.e., *Per1/2* null mice) fed either ad libitum or exclusively during the night. We found that a similar fraction of lipids (~17%) oscillated in both mouse strains, most notably triglycerides (TAG), but with completely different phases. Moreover, several master lipid regulators and enzymes involved in TAG metabolism retained their circadian expression in clock-disrupted mice. Feeding time had a prominent effect on the phase and levels of hepatic TAG in both WT and *Per1/2* null mice. Remarkably, upon nighttime-restricted feeding, WT mice exhibited a sharp decrease (~50%) in hepatic TAG levels. Our

findings suggest that circadian clocks and feeding time dictate the phase and levels of hepatic TAG accumulation; however, oscillations in TAG can persist in the absence of a functional clock.

RESULTS

Mouse Liver Lipidome

To obtain a temporal depiction of the circadian changes in lipid abundance in mouse liver, we performed a wide lipidomic analysis. WT mice were fed ad libitum and housed under a 12 hr light/dark regimen for several consecutive days. Throughout the last day of the experiment, animals were maintained in constant darkness and sacrificed every 4 hr. Livers were harvested, and lipids were identified and quantified by shotgun lipidomics (Han et al., 2012). Altogether 159 lipids were measured (see Table S1A available online). These include different triglycerides (TAG), phospholipids (i.e., phosphatidylinositol [PI], phosphatidylcholine [PC], lysophosphatidylcholine [LPC], phosphatidylethanolamine [PE], phosphatidylserine [PS], phosphatidic acid [PA], and phosphatidylglycerol [PG]), sphingolipids (i.e., ceramide [CER] and sphingomyelin [SM]), cardiolipin (CL), and acyl carnitine (CAR). A nonparametric algorithm, JTK_CYCLE (Hughes et al., 2010), was used to identify lipids that display circadian rhythmicity. Out of the 159 lipids, 27 (~17%) were identified as oscillating with p value < 0.05 (Figures 1A and S1; Table S1B). Among the 27 oscillating lipids, the majority consisted of TAG species (~33%), (Figures 1B and 1C). When dissected

based on their types, it appeared that oscillating lipid species were highly abundant among PI (46%) and enriched among TAG (26%) and PC (26%) (Figure 1D). Analysis of the phase distribution of the different oscillating lipids revealed that the vast majority of them peaked during the subjective light phase. Namely, 23 out of the 27 reached their zenith levels between circadian time (CT) 1 and CT8 (Figure 1E). Hence, we concluded that oscillating lipids mostly consist of TAG species and are predominantly temporally gated to reach their zenith levels during the subjective light phase.

Circadian Expression of Enzymes Participating in Hepatic TAG Metabolism

Hepatocytes harbor the ability to synthesize, store, and catabolize TAG. A prominent fraction (i.e., 33%) of oscillating lipids consisted of TAG species (Figure 1B). Strikingly, all oscillating TAG peaked around CT8 and dipped around CT20 with amplitude of about 2-fold (Figures 1C and 1E), suggesting that their accumulation in the liver is temporally controlled. To corroborate these findings, we examined the circadian expression profile of multiple enzymes that participate in TAG metabolism in WT mice. The glycerol-3-phosphate pathway is the predominant biosynthesis pathway for TAG in the liver and consists of several successive steps (Figure 2A) (Takeuchi and Reue, 2009). The first key and rate-limiting step is the acylation of glycerol 3-phosphate by glycerol-3-phosphate acyltransferase (GPAT) enzymes to synthesize lysophosphatidic acid (LPA). Subsequently, an additional fatty acid is transferred to LPA by the family of 1-acylglycerol-3-phosphate acyltransferase (AGPAT) enzymes to produce phosphatidate (PA). Diacylglycerol (DAG) is synthesized from PA by the Lipin family of proteins, through their phosphatidate phosphatase-1 (PAP) enzymatic activity. Finally, DAG is converted to TAG through the action of diacylglycerol acyltransferase (DGAT) enzymes.

WT mice were sacrificed under constant darkness at 4 hr intervals throughout the day, livers were harvested, and RNA was prepared. The expression level of the different enzymes was quantified by real-time PCR. Remarkably, we identified circadian accumulation pattern for multiple enzymes in the glycerol-3-phosphate pathway, covering all subsequent steps involved in TAG biosynthesis (Figure 2B). While *Gpat1* mRNA levels were relatively constant throughout the day, *Gpat2* exhibited shallow circadian oscillations with zenith levels around CT0 and nadir levels around CT12. Next, both *Agpat1* and *Agpat2* transcripts accumulated in a circadian manner, albeit with a phase difference: the former peaked at CT16, while the latter peaked at CT0. Subsequently, the mRNA expression profile of both *Lpin1* and *Lpin2* was circadian, with zenith levels at CT12 and nadir levels at CT0. Finally, the transcript levels of *Dgat2* but not *Dgat1* were oscillating with a phase similar to that of *Lpin1* and *Lpin2*. These findings demonstrated that the glycerol-3-phosphate pathway is expressed in a circadian manner.

Hepatic TAG catabolism occurs through the activity of multiple lipases, releasing one fatty acid at a time, producing diacylglycerols and eventually glycerol (Figure 2A) (Quiroga and Lehner, 2012). To examine the circadian regulation of TAG catabolism in the liver, we analyzed the expression levels of two major hepatic lipases, the patatin-like phospholipase domain containing 3 (*Pnpla3*) and the lysosomal acid lipase (*Lipa*) (Quiroga and

Lehner, 2012). The mRNA expression levels of both *Pnpla3* and, to a much lesser extent, *Lipa* were circadian, with nadir levels around CT12 (Figure 2B). Taken together, our data evinced that multiple hepatic enzymes that participate in TAG homeostasis oscillate in a circadian manner. Conceivably, these enzymes function in concert to generate the above-described circadian oscillation of TAG in the liver.

Liver Lipidome Analysis of *Per 1/2* Null Mice

To address the clock dependency of the above-illustrated circadian profile of lipid accumulation in mouse liver, we performed a lipidomic analysis on liver samples obtained from *Per1/2*^{-/-} mice. Mice lacking both PER1 and PER2 exhibit arrhythmic locomotor activity under constant darkness, and circadian expression of core clock and clock-controlled genes is largely diminished (Zheng et al., 2001). Consistently, expression levels of core clock genes (i.e., *Bmal1*, *Clock*, *Cry*, and *Rev-erb α*) were relatively constant throughout the day, and as expected, *Per1* and *Per2* mRNA levels were undetectable in these mice (Figure S2).

Per1/2 null mice were fed ad libitum and sacrificed under constant darkness at 4 hr intervals throughout the day. Livers were harvested and lipids were identified and quantified by shotgun lipidomics (Table S2A). Analysis based on the JTK_CYCLE algorithm evinced that out of the 159 classified and quantified lipids, 25 (~16%) were identified as oscillating with $p < 0.05$ (Figures 3A and S3; Table S2B). The preponderance of them consisted of TAG (~76%), (Figures 3B, 3C, and S3). Moreover, when dissected based on their types, oscillating lipid species were highly enriched among the TAG (~54%), (Figure 3D). Phase distribution analysis revealed a mirror image to that obtained for WT samples (compare Figure 1E with Figure 3E). Thus, all oscillating TAG reached their peak levels during the late night, more specifically between CT20 and CT0 (Figures 3C, 3E, and S3). The observed oscillations in lipid accumulation, most notably TAG in the absence of PER1/2, suggest that their circadian accumulation might be clock independent.

The Effect of Feeding Time on Circadian TAG Accumulation in the Liver

Food is a major source for lipids in general and triglycerides in particular. To examine the direct effect of food ingestion on the accumulation pattern of TAG in the liver, we performed a limited lipidomic analysis, this time measuring only TAG species in WT and *Per1/2* null mice, fed exclusively during the night, namely between CT12 and CT0 (Tables S3A and S4A). JTK_CYCLE based analysis showed that out of 35 measured TAG species, 16 were oscillating in WT and 18 in *Per1/2* null mice (~46 and ~51%, respectively) ($p < 0.05$) (Figure S4; Tables S3B and S4B). All oscillating TAG species peaked around CT12 in both mouse strains (Figure S4). However, while WT mice exhibited a gradual buildup and decline of TAG with zenith levels at CT12, *Per1/2*^{-/-} mice displayed a prominent and sharp peak in TAG accumulation at CT12 (Figures 4A and S4). The surge in TAG accumulation in night-fed animals was probably due to lack of food ingestion prior to CT12, as mice were sacrificed shortly before food was provided. The prominent phase shift in TAG accumulation from around CT20 in *Per1/2*^{-/-} mice and CT8 in WT mice fed ad

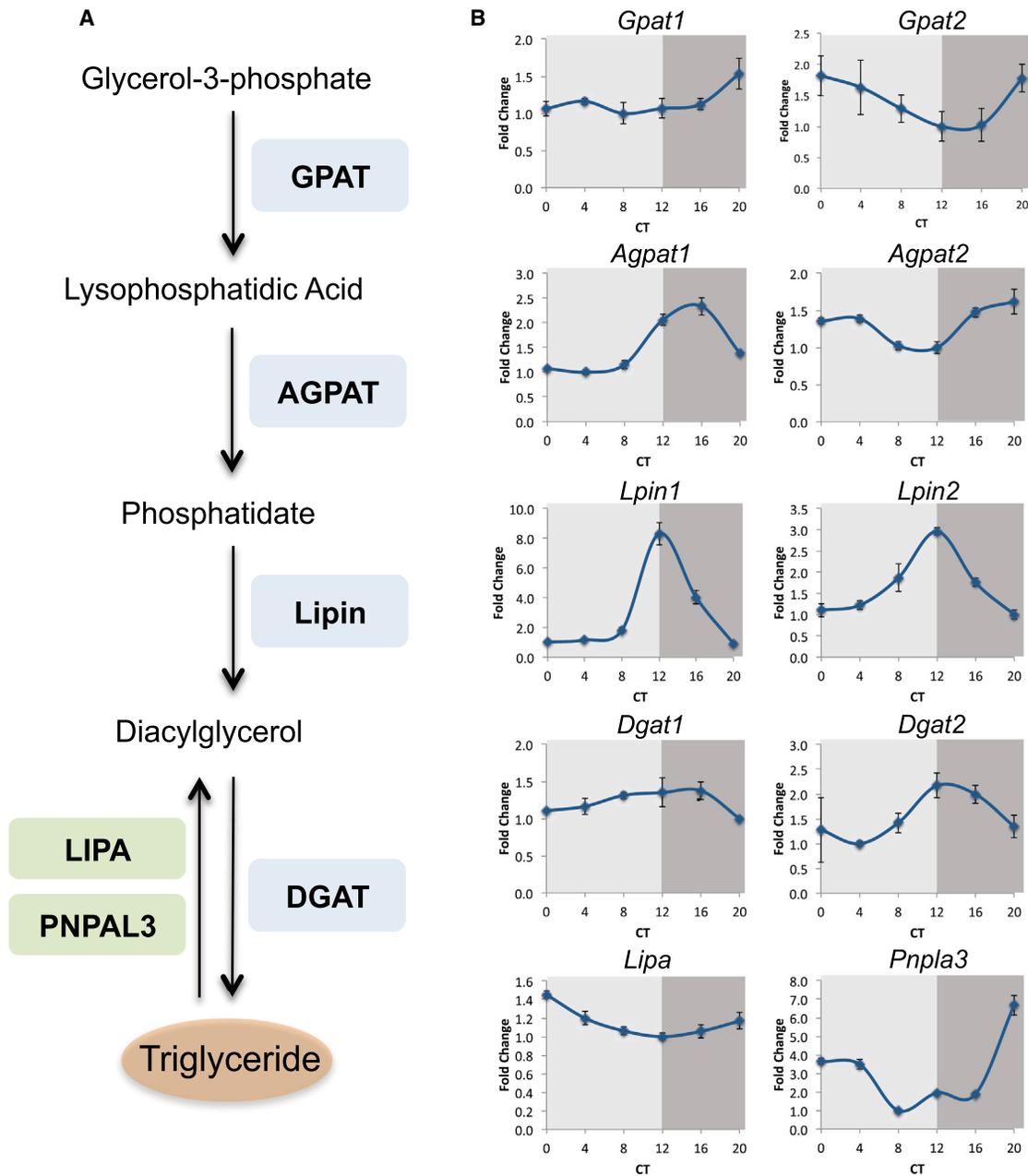


Figure 2. Circadian Regulation of Hepatic Triglyceride Metabolism

(A) Schematic depiction of the triglyceride biosynthesis pathway (glycerol-3-phosphate pathway) and triglyceride catabolism. Glycerol-3-phosphate acyltransferase (GPAT), 1-acylglycerol-3-phosphate acyltransferase (AGPAT), diacylglycerol acyltransferase (DGAT), patatin-like phospholipase domain-containing 3 (PNPLA3), and lysosomal acid lipase (LIPA).

(B) WT mice were sacrificed under constant darkness at 4 hr intervals throughout the day. Total RNA was prepared from liver, and mRNA expression levels were determined by quantitative real-time PCR and presented as fold change relative to the lowest value. Data are presented as mean \pm SD, with a mix of four animals per time point. CT, circadian time.

libitum to CT12 upon night feeding implied that feeding-fasting cycles can shape the phase of TAG accumulation in mouse liver.

Analysis of Feeding and Clock Dependency of Circadian Hepatic TAG Accumulation

Hitherto, we obtained a temporal depiction of circadian hepatic TAG accumulation in WT and *Per1/2*^{-/-} mice, fed either ad libitum

or exclusively during the night (Figure 4A). Altogether, we identified circadian oscillations in 9, 16, 19, and 18 out of 35 quantified TAG species, which peaked at ~ZT8, ~ZT12, ~ZT20, and ~ZT12 in WT fed ad libitum, WT night-fed, *Per1/2*^{-/-} fed ad libitum, and *Per1/2*^{-/-} night-fed mice, respectively. By mass this corresponded to 22%, 41%, 61%, and 53% of total TAG liver mass, respectively. Hence, these changes were also reflected to some extent in

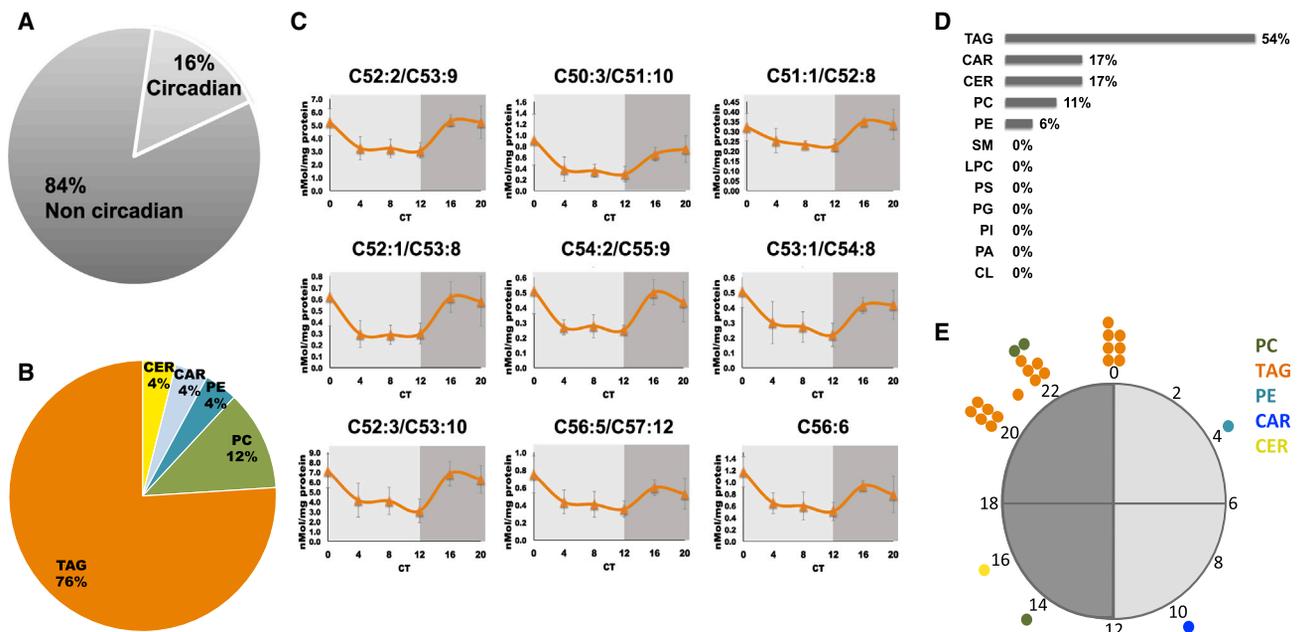


Figure 3. Liver Lipidome Analysis of *Per 1/2* Null Mice

- (A) The percentage of lipids that were found to exhibit a circadian pattern of accumulation in livers of *PER1/2*^{-/-} mice based on JTK_CYCLE analysis (six time points, n = 4 for each, p < 0.05). Out of the 159 measured and identified lipids, 25 lipids exhibited circadian pattern of accumulation.
- (B) Oscillating lipids species distributed according to their types.
- (C) Accumulation profiles of representative oscillating TAG presented as mean ± SD. CT, circadian time.
- (D) The percentage of the oscillating lipid species within each lipid type.
- (E) Daytime distribution of peak phases of oscillating lipids. TAG is in orange, PC in green, PE in light blue, CAR in blue, and CER in yellow. See also Figures S2 and S3.

circadian oscillations of total hepatic TAG levels, particularly in the case of WT night-fed, *Per1/2*^{-/-} fed ad libitum, and *Per1/2*^{-/-} night-fed mice (p < 0.05 based on the JTK_CYCLE analysis) (Figure 4B).

The ability of feeding time to shift the phase of TAG accumulation raised the question whether the striking difference in the phase of TAG oscillations in WT versus *Per1/2*^{-/-} mice fed ad libitum might be the outcome of different daily eating habits of the two genotypes. Therefore, we temporally monitored the food intake of WT and *Per1/2* null mice. WT mice ingested ~70% of their total daily food intake during the subjective dark phase and ~30% during the subjective light phase, whereas *Per1/2*^{-/-} mice largely differed in their eating habits and ingested relatively equal amounts of food throughout the day (Figures 4C and S5). Altogether, both WT and *Per1/2*^{-/-} consumed a similar amount of food throughout the day (Figure 6C). Thus, though *Per1/2* null mice differed in their eating habits from WT mice, their eating behavior could not fully explain the circadian accumulation pattern of TAG in these mice, as they consumed food constantly throughout the day.

Dissection of Oscillating TAG Species Based on Feeding Time and Circadian Clocks

To uncover principles that shape TAG landscape in the liver, namely to identify which TAG species are primarily clock driven and which are mainly regulated by food intake, we dissected the overlap between the different TAG species that oscillated under the four different experimental setups (i.e., WT fed ad libitum, WT

night fed, *Per1/2*^{-/-} fed ad libitum, and *Per1/2*^{-/-} night fed), (Figure 5). There was a relatively small commonality between the oscillating TAG species in WT mice fed ad libitum and *Per1/2*^{-/-} mice fed ad libitum (four out of nine and 19, respectively), (Figure 5A). Upon nighttime feeding, the number of oscillating TAG species in WT mice increased from 9 to 16. All nine species that oscillated under ad libitum regimen continued to oscillate in night-fed animals (Figure 5B). In sharp contrast, there was a little overlap between the oscillating TAG species under ad libitum and nighttime feeding in *Per1/2*^{-/-} mice. Only six species overlapped out of the 19 that oscillated in ad libitum and 18 in night-fed animals (Figure 5C). Remarkably, we observed an extensive overlap between the TAG species that oscillate in WT night-fed and *Per1/2*^{-/-} night-fed animals (15 out of 16 and 18, respectively) (Figure 5E). These findings suggest that there is a distinct population of oscillating TAG species that respond to feeding-fasting cycles independently of PER1/2 (Figure S6A). Strikingly, the TAG species that oscillated in *Per1/2*^{-/-} fed ad libitum largely differed from the one that appeared to be food driven (Figure S6B). This in line with our findings that *Per1/2*^{-/-} mice fed ad libitum eat constantly throughout the day and hence that feeding-fasting cycles do not determine their circadian TAG accumulation.

To further characterize these different TAG populations, we plotted them according to the number of carbons. TAG consist of a glycerol backbone, to which three molecules of fatty acids are attached. Hence the carbon number provides insight regarding the fatty acid composition of the different TAG. In

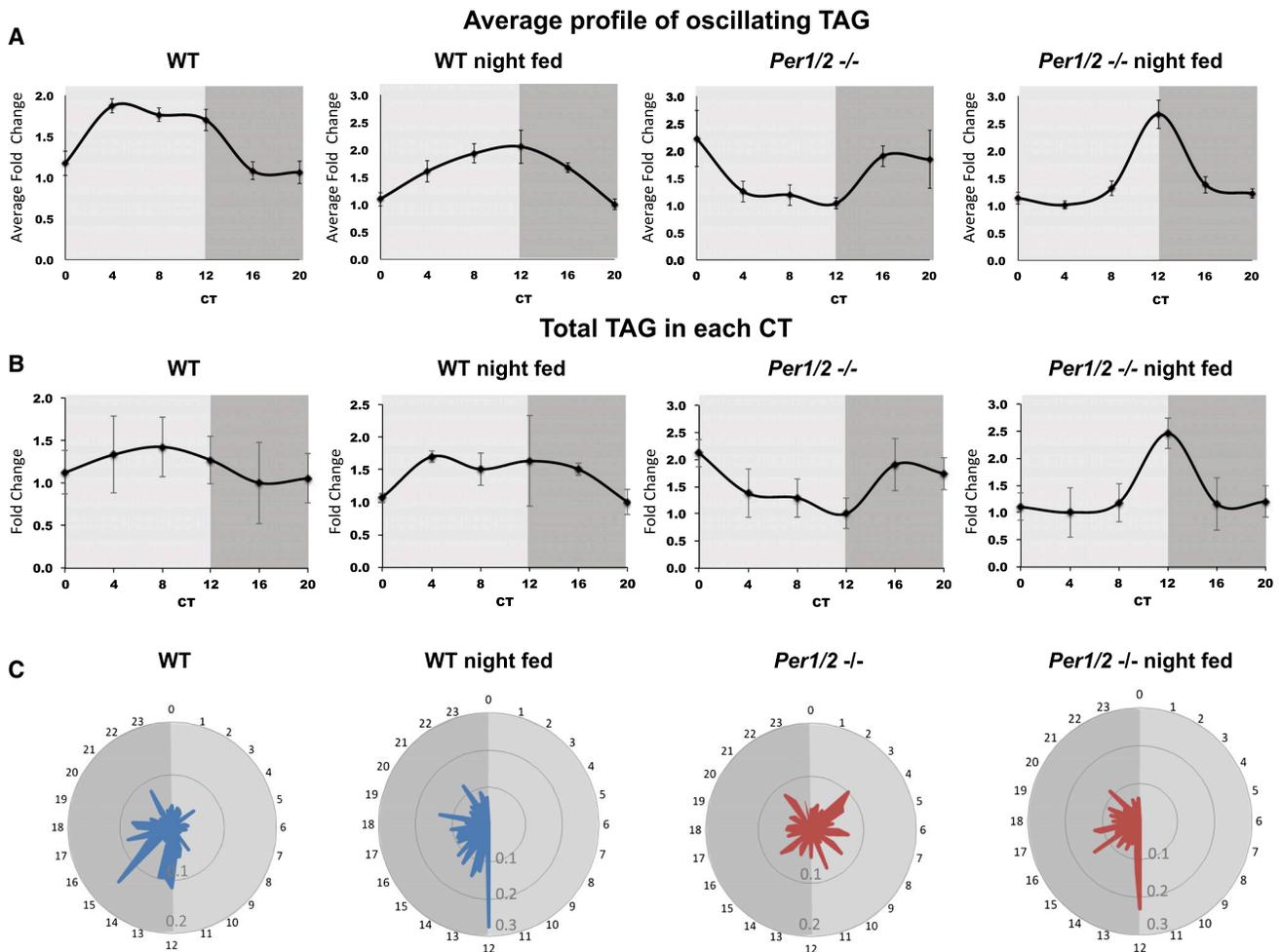


Figure 4. Analysis of Feeding and Clock Dependency of Circadian Hepatic TAG Accumulation

(A) Profile of oscillating TAG identified based on JTK_CYCLE analysis for WT and *Per1/2*^{-/-} mice fed either ad libitum or exclusively during the night. Data are presented as mean ± SD of fold induction for all oscillating TAG relative to the lowest point for each one.

(B) Accumulation profiles of total TAG levels in livers of WT and *Per1/2*^{-/-} mice fed either ad libitum or exclusively during the night. Data are presented as mean ± SD of fold induction for total TAG levels relative to the lowest point. The corresponding p values based on JTK_CYCLE analysis were 0.31, 0.03, 0.02, and 0.02, respectively.

(C) A radar plot presenting the circadian food consumption of WT and *Per1/2*^{-/-} mice fed either ad libitum or exclusively during the night. The time of day is indicated in hours, and the y axis shows the food consumption in grams. See also Figures S4, S5, and S7 and Table S6.

WT mice fed ad libitum, the oscillating TAG was comprised of 54 and higher number of carbons, indicating that they primarily contain long-chain fatty acids (C14-20), whereas in *Per1/2*^{-/-} mice fed ad libitum a large fraction included also TAG with carbon number below 54, suggesting that they also contain medium-chain fatty acid (C8-12) (Figure S6C). Under night-fed conditions, both mice strains exhibited a very similar distribution (Figure S6D), which actually reflects the strong overlap between the oscillating species in these two groups (Figure 5E). Overall, these analyses suggested that the oscillating TAG species in *Per1/2*^{-/-} fed ad libitum largely differ from the TAG identified in all other tested conditions (i.e., WT fed at ad libitum, WT night fed, and *Per1/2*^{-/-} night fed). Taken together with our finding that *Per1/2*^{-/-} mice ingest food constantly throughout the day, it is unlikely that the observed TAG oscillations in *Per1/2*^{-/-} are food driven.

The Effect of Feeding Time and Circadian Clocks on Total TAG Levels in the Liver

Previous experiments performed with *Per2* single knockout mice have demonstrated that their daily food intake is identical to WT mice but that their total TAG plasma levels are drastically reduced (~50%) (Grimaldi et al., 2010). In line with these observations, quantification of the total hepatic TAG levels throughout the day revealed a sharp decrease (~50%) in hepatic TAG content of *Per1/2*^{-/-} mice compared with WT mice (Figure 6A). The level of none of the other measured lipid types was decreased in *Per1/2* null mice compared with WT mice. The levels of PE, PG, SM, PC, LPC, and CAR did not differ between the two mouse strains, whereas CER, PI, CL, PS, and PA levels in *Per1/2* null mice were elevated (Figure 6A). Interestingly, 2 weeks of nighttime feeding drastically reduced the total hepatic TAG content in WT mice (~50% reduction) (Figure 6B). By contrast, nighttime

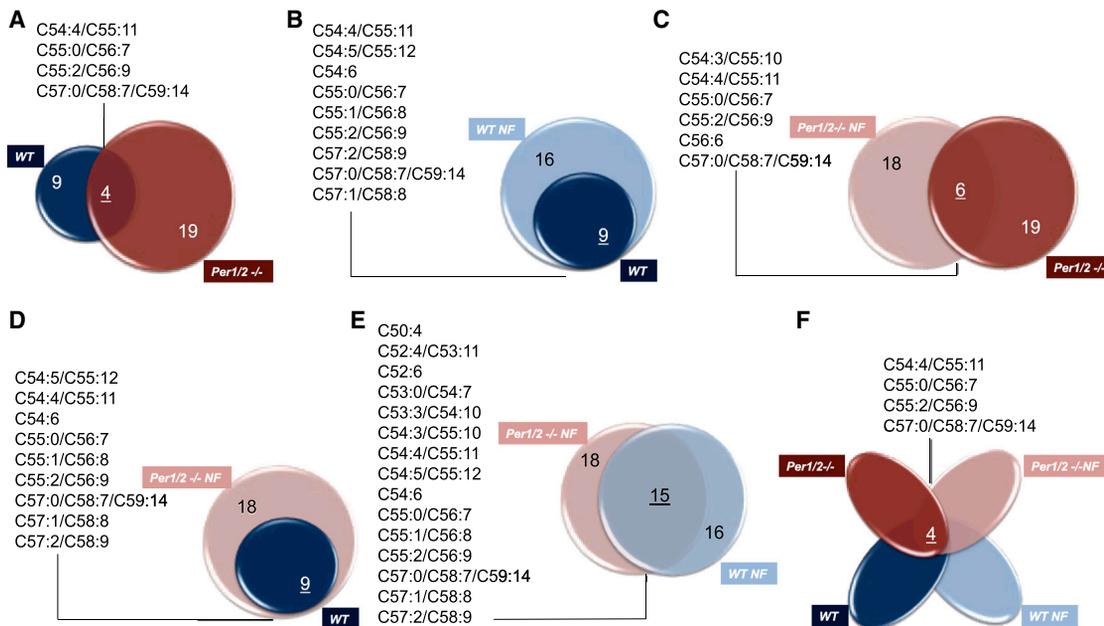


Figure 5. Dissection of Oscillating TAG Species Based on Feeding Time and Circadian Clocks

Overlap analysis between the specific TAG species that oscillated under the different experimental setup.

(A) WT mice fed ad libitum and *Per1/2*^{-/-} mice fed ad libitum.

(B) WT night-fed mice and WT mice fed ad libitum.

(C) *Per1/2*^{-/-} night-fed mice and *Per1/2*^{-/-} mice fed ad libitum.

(D) WT mice fed ad libitum and *Per1/2*^{-/-} mice night fed.

(E) WT night-fed mice and *Per1/2*^{-/-} night-fed mice.

(F) WT mice fed ad libitum, WT night-fed mice, *Per1/2*^{-/-} mice fed ad libitum, and *Per1/2*^{-/-} night-fed mice. For each scheme, the list of overlapping TAG species is detailed. NF, night fed.

See also Figure S6.

feeding of *Per1/2*^{-/-} mice resulted in a significant increase of about 25% in hepatic TAG content (Figure 6B). This indicated that time-restricted feeding drastically affects the TAG levels in the liver. These differences were evident in almost every single TAG species that was quantified in our lipidomic analysis (Table S5). It should be noted that under all four tested conditions (i.e., WT mice fed ad libitum or night fed, *Per1/2*^{-/-} mice fed ad libitum or night fed), we did not observe any significant differences in the total daily food intake of the mice (Figure 6C), indicating that the observed changes in hepatic TAG levels were not due to differences in their food consumption. These findings implied that both feeding and clock-dependent mechanisms not only dictate the phase of hepatic TAG accumulation but also play a prominent role in determining the total hepatic TAG levels throughout the day.

Feeding- and Clock-Driven Expression of Enzymes and Master Regulators of Hepatic TAG Metabolism

To further corroborate these findings and dissect the clock and/or feeding dependency, we determined the expression profile of enzymes participating in TAG homeostasis in WT and *Per1/2* null mice fed either ad libitum or exclusively during the night (Figure 7A). Comparison under ad libitum feeding conditions demonstrated that the expression profiles of several enzymes in TAG metabolism, notably *Gpat1*, *Gpat2*, *Agpat2*, *Lpin1*, *Lpin2*, *Dgat2*, *Lipa*, and *Pnpla3*, differ between the two genotypes.

Both *Lpin1* and *Gpat2* mRNA levels oscillated in WT mice and were relatively low and constant in *Per1/2* null mice, indicating that they are expressed in a clock-dependent manner. Remarkably, several enzymes exhibited circadian oscillations in *Per1/2* null mice (i.e., *Gpat1*, *Agpat1*, *Agpat2*, *Lpin2*, *Lipa*, and *Pnpla3*). Notably, although *Pnpla3* mRNA levels were sharply reduced in *Per1/2* null mice compared to WT mice, *Pnpla3* continued to oscillate in *Per1/2*^{-/-} mice with ~5-fold amplitude. The expression profile of *Gpat1*, *Agpat2*, *Lpin2*, *Lipa*, and *Pnpla3* was phase shifted in *Per1/2*^{-/-} mice. For example, *Agpat2* and *Pnpla3* mRNA levels accumulated around CT0 in WT mice and around CT16 in *Per1/2*^{-/-} mice. Similarly, *Lpin2* accumulated about 8 hr earlier in *Per1/2*^{-/-} mice (i.e., ~CT4 compared to ~CT12). The observed oscillations in the absence of PER1/2 suggest that the circadian accumulation of these enzymes might be clock independent.

To examine the direct effect of feeding time, we compared the expression pattern in mice fed either ad libitum or exclusively during the night. The circadian mRNA expression pattern of *Lpin1* and *Gpat2* persisted under both feeding conditions, suggesting that their accumulation is primarily clock dependent and less affected by feeding time. Altogether, in WT mice, nighttime feeding had very little effect on the phase of oscillating enzymes. By contrast, in *Per1/2*^{-/-} the expression phase of *Gpat1*, *Agpat2*, *Lpin2*, *Lipa*, and *Pnpla3* was altered upon nighttime feeding, pointing out that their circadian expression responds

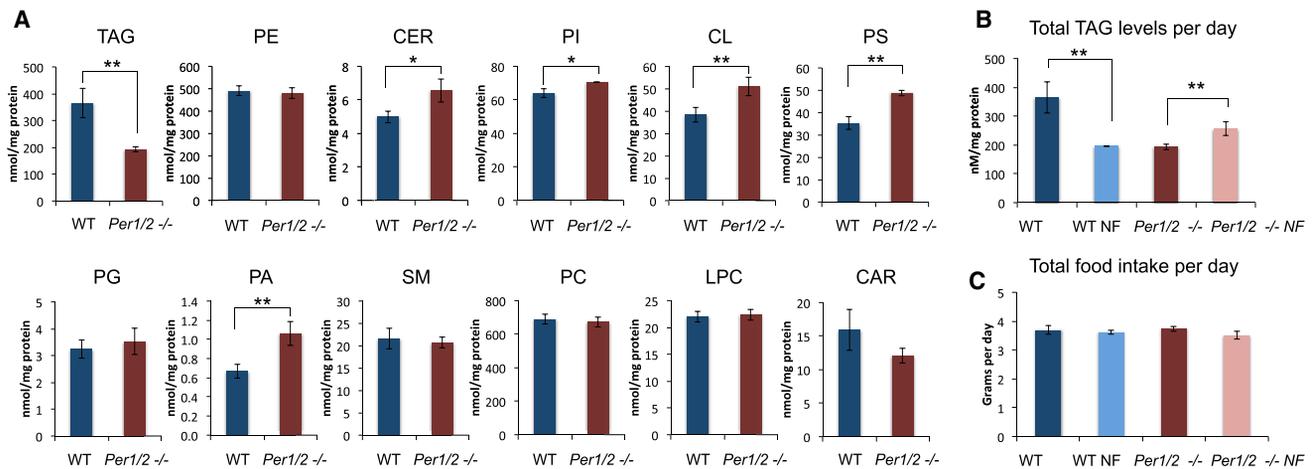


Figure 6. The Effect of Feeding Time and Circadian Clocks on Total TAG Levels in the Liver

(A) Comparison of the total hepatic levels of the different lipid types, quantified throughout the day, between WT and *Per1/2*^{-/-} mice fed ad libitum.

(B) Comparison of the total hepatic TAG levels, quantified throughout the day in WT mice fed ad libitum, WT night-fed mice, *Per1/2*^{-/-} mice fed ad libitum, and *Per1/2*^{-/-} night-fed mice.

(C) Comparison of the total daily food intake of WT mice fed ad libitum, WT night-fed mice, *Per1/2*^{-/-} mice fed ad libitum, and *Per1/2*^{-/-} night-fed mice. The data are presented on a bar graph (mean ± SEM, n = 4 for the lipid analysis and n = 8 for the food consumption). NF, night fed. *p < 0.05; **p < 0.01.

to changes in feeding time. Overall, our analysis evinced that the circadian expression of several hepatic enzymes participating in TAG metabolism is clock dependent, whereas other enzymes retain their circadian expression in the absence of a functional clock and respond to feeding.

The circadian oscillations in expression of enzymes participating in TAG homeostasis in *Per1/2*^{-/-} mice prompted us to examine the expression profile of master regulators of lipid metabolism in these mice. The nuclear receptors, peroxisome proliferator-activated receptor alpha (PPAR α) and gamma (PPAR γ), that promote the expression of genes involved in fatty acids uptake, utilization, and catabolism are expressed in a circadian manner (Yang et al., 2006). Indeed, both PPAR α and PPAR γ mRNA levels oscillated in livers of WT mice; the former peaked around CT8 and the latter around CT4 (Figure 7B). Remarkably, the circadian expression pattern of PPAR α persisted in *Per1/2* null mice with a similar magnitude and amplitude. A 4-fold reduction in the mRNA levels of PPAR γ was observed in *Per1/2* null mice. Yet, the PPAR γ transcript levels retained their circadian expression and peaked around CT8 (~2-fold induction). Hence, both PPAR α , and PPAR γ cycled in the absence of PER1/2 proteins. The sterol regulatory element-binding protein (SREBP1), a transcription factor that regulates the expression of genes involved in cholesterol and lipid metabolism, accumulates in a daily manner (Le Martelot et al., 2009). As expected, in WT mice, *SREBP1c* mRNA levels oscillated in a circadian manner with zenith levels at ~CT16 and nadir levels at ~CT4. By contrast, the expression levels of *SREBP1c* in *Per1/2* null mice were relatively low and constant throughout the day (Figure 7B). The expression levels of other central regulators of lipid homeostasis, such as hepatocyte nuclear factor 4 (HNF4) and liver X receptors alpha (LXR α) and beta (LXR β), were relatively similar and constant throughout the day in both mouse strains (data not shown).

Thus, we concluded that though the core clock circuitry is non-oscillating in the absence of PER1/2 (Figure S2), we do observe

circadian oscillations in the expression of several master lipid regulators and enzymes that participate in TAG homeostasis.

DISCUSSION

We provide a comprehensive temporal and quantitative analysis of lipid accumulation in livers of WT and clock-disrupted mice fed either ad libitum or exclusively during the night. In view of the wide involvement of lipids in various key cellular functions (e.g., energy storage and provision, membrane composition, and signal transduction), our finding that ~17% of all measured lipids in mouse liver exhibit circadian pattern of accumulation has wide ramifications. Remarkably, our liver lipidomic analysis showed that numerous TAG species accumulate in a circadian manner, all reaching their peak levels around CT8 (Figure 1). In this conjuncture, expression analysis of enzymes participating in TAG metabolism in the liver revealed a circadian accumulation pattern for multiple enzymes covering all subsequent steps involved in TAG biosynthesis (Figure 2).

As previously mentioned, circadian transcriptome analyses have been highly valuable in identifying core clock and output genes, including many genes involved in various metabolic pathways. Nevertheless, since the majority of cellular processes are tightly regulated at multiple levels beyond transcription (e.g., translation rate, posttranslational modification, and protein stability) these approaches are limited and provide only a partial depiction. This is particularly pertinent for metabolic pathways that often rely on intricate cascade of enzymatic reactions and are modulated by enzyme activity and substrate availability. Hence, measurements of the absolute levels of metabolites are essential. Indeed, multiple enzymes involved in TAG metabolism in the liver were expressed in a circadian manner (Figure 2), yet they exhibited a wide range of expression phases. Thus, solely based on the mRNA expression data, it would have been difficult to predict whether the final products (i.e., TAG) would

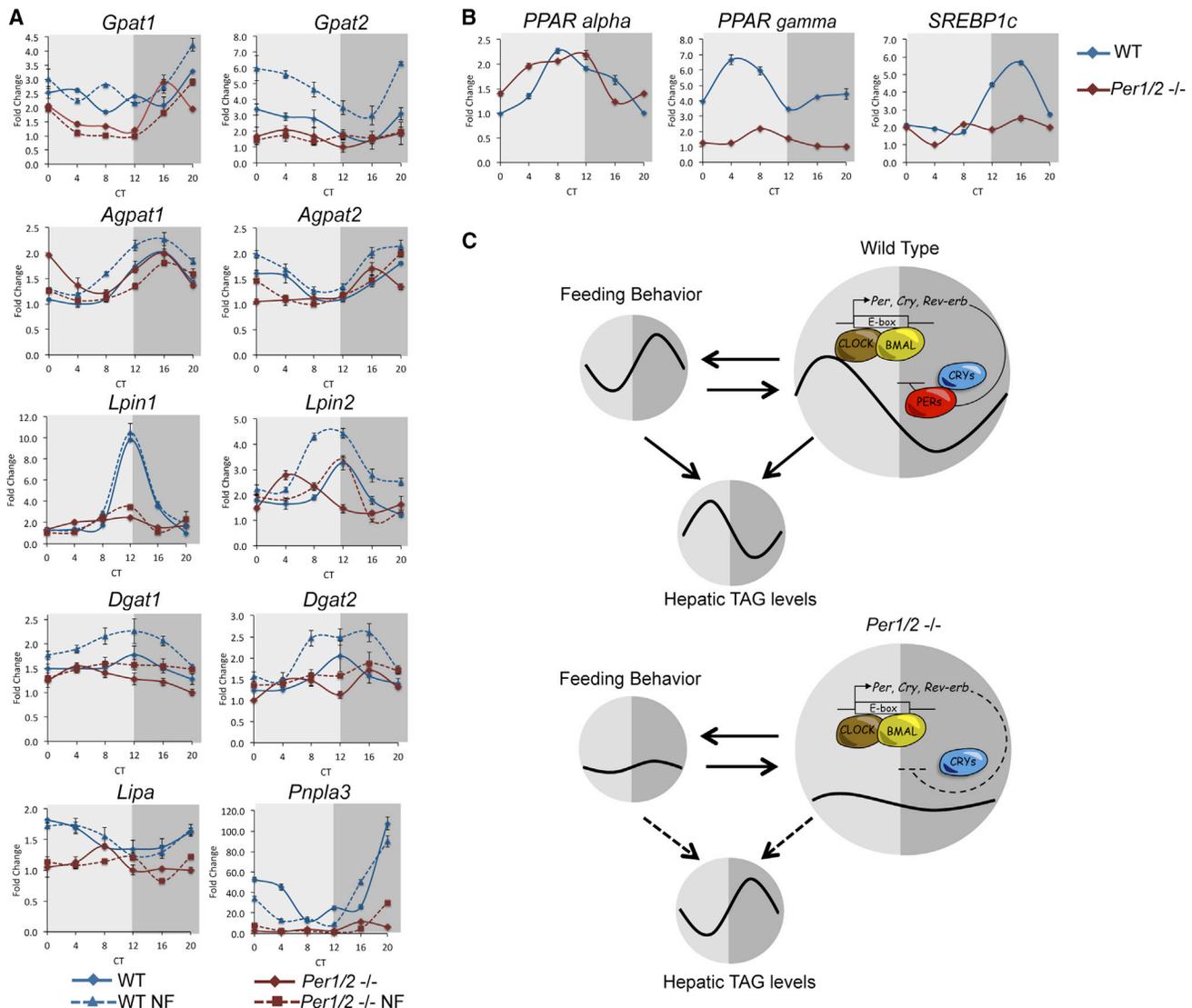


Figure 7. Circadian Expression of Enzymes Participating in TAG Homeostasis and Master Lipid Regulators in WT and *Per1/2* Null Mice

(A) WT and *Per1/2*^{-/-} mice, fed either ad libitum or exclusively during the night, were sacrificed under constant darkness at 4 hr intervals throughout the day. Total RNA was prepared from liver, and mRNA expression levels of enzymes participating in TAG metabolism were determined by quantitative real-time PCR and presented as fold change relative to the lowest value.

(B) Circadian mRNA expression profiles of master lipid regulators in WT and *Per1/2*^{-/-} mice fed ad libitum. Data are presented as mean \pm SD, with a mix of four animals per time point. Dark gray represents the subjective night and light gray the subjective day.

(C) A schematic model depicting the intricate circadian regulation of hepatic TAG levels by circadian clocks and feeding behavior. NF, night fed; CT, circadian time.

accumulate in a circadian manner and identify their actual phase. Thus, our analysis clearly showed that in contrast to the disparate expression of enzymes involved in TAG homeostasis, all oscillating TAG peak around CT8 in WT mice fed ad libitum (Figure 1). A plausible explanation for that is that posttranscriptional control of these enzymes in a temporal manner might serve as a determinant of TAG synthesis. Several examples in the literature highlight the role of posttranscriptional control of enzymes in TAG biosynthesis. Both AMP-activated protein kinase (AMPK), a sensor for cellular energy levels, and casein kinase have been implicated in the phosphorylation and regulation of GPAT enzymatic activity (Coleman and Lee, 2004). Notably, AMPK and casein kinase are significant components within the core

clock circuitry (Dibner et al., 2010). Phosphorylation of Lipin1, on multiple serine and threonine residues, in response to various stimuli (e.g., insulin and amino acids) regulates its subcellular localization and its PAP enzymatic activity (Harris et al., 2007; Peterson et al., 2011). Lastly, posttranscriptional control via translation regulation has been also linked to DGAT accumulation (Coleman and Lee, 2004).

Previous genome-wide analyses of the temporal DNA binding of BMAL1 and additional core clock proteins using chromatin immunoprecipitation combined with deep sequencing showed that their binding sites are enriched among genes involved in lipid metabolism, notably triglyceride biosynthesis (Cho et al., 2012; Koike et al., 2012; Rey et al., 2011). Binding sites for

BMAL1, CLOCK, PER1/2, and CRY1/2 were identified for multiple enzymes in the glycerol-3-phosphate pathway. Specifically, BMAL1, CLOCK, PER1, and CRY1/2 were found to bind the same intergenic loci, upstream to the *Lpin1* transcription-starting site, albeit with a different phase. Similarly, BMAL1, CLOCK, PER1/2, and CRY1/2 were present on the same DNA region, upstream to the *Gpat2* transcription-starting site, though during different times of the day (~ZT4 for BMAL/CLOCK and ~ZT16 for PER/CRY). Hence, it is conceivable that both *Lpin1* and *Gpat2* are direct targets of BMAL/CLOCK, PER/CRY transcriptional regulation, as the prominent mRNA circadian expression of both *Lpin1* and *Gpat2* was strongly dampened in clock-disrupted mice (Figure 7).

Interestingly, we found that a similar fraction of lipids (~17%) were oscillating in both WT and *Per1/2* null mice fed ad libitum, most notably TAG. However, they largely differed in their accumulation phase and composition. These observations are intriguing, as mice lacking both PER1 and PER2 are arrhythmic under constant darkness and their circadian expression of core clock genes is largely abolished (Figure S2) (Zheng et al., 2001). This raises the question of what the molecular mechanisms are that drive the circadian oscillations in TAG accumulation in the absence of a functional clock. We do show that feeding-fasting cycles can strongly shape the phase of TAG accumulation in mouse liver and that *Per1/2*^{-/-} mice differ in their eating habits compared to WT mice. However, their feeding behavior cannot fully explain the oscillations of TAG in these mice, as they consume equal amounts of food throughout the day, and hence food ingestion cannot serve as a timing cue. Moreover, overlap analyses of the oscillating TAG species, identified in WT and *Per1/2*^{-/-} mice fed ad libitum or exclusively during the night, point out that the TAG species that oscillated in *Per1/2*^{-/-} mice fed ad libitum largely differed from the oscillating TAG species found in night-fed *Per1/2*^{-/-} mice and WT mice (Figure 5). Conceivably, the TAG species that oscillated in *Per1/2*^{-/-} mice fed ad libitum are comprised of a distinct population that is not driven by feeding-fasting cycles.

Further studies are required to identify the molecular mechanisms that drive the circadian accumulation of TAG in the absence of a functional clock. It should be noted that while the expression of core clock genes is relatively constant throughout the day in *Per1/2* null mice (Figure S2), and these mice exhibit arrhythmic locomotor activity and feeding behavior under constant darkness (Figure 4C) (Zheng et al., 2001), our analyses do demonstrate the persistence of circadian oscillation in mRNA expression of enzymes that participate in TAG metabolism and master lipid regulators such as PPARs (Figure 7). Hence, this suggests that different circadian outputs (e.g., locomotor activity, feeding behavior, TAG accumulation, mRNA expression) are selectively affected in the absence of PER1/2; some do persist, while others dampen.

Recently, it has been shown that time-restricted feeding for more than 3 months can prevent obesity, hepatic steatosis, and metabolic syndrome in mice fed a high-fat diet (Hatori et al., 2012; Sherman et al., 2012). Time-restricted feeding primarily protected mice from the adverse effects of a high-fat diet, but several metabolic parameters were also significantly improved upon time-restricted feeding of standard diet, among them hepatic steatosis (Hatori et al., 2012; Sherman et al.,

2012). In this conjuncture, our analysis evinced that time-restricted feeding of regular chow diet in WT mice for a time period as short as 2 weeks results in a dramatic decrease (~50%) in total TAG levels in the liver, whereas the total food intake was unchanged. Elevated plasma and hepatic TAG levels are associated with hepatic steatosis and liver malfunction. The striking effect of nighttime feeding on hepatic TAG levels is of obvious clinical importance, as current pharmacological interventions are less efficient and often associated with various adverse side effects (McKenney and Sica, 2007).

In summary, our study evinces that both circadian clocks and feeding-fasting cycles play a prominent role in the regulation of circadian TAG accumulation and total TAG levels in the liver. Nevertheless, in the absence of a functional clock (i.e., *Per1/2*^{-/-} mice) and thus lack of feeding rhythms, circadian oscillations in hepatic TAG levels do persist, albeit with a complete different phase. This suggests that additional mechanisms play a role in their circadian accumulation (Figure 7C).

EXPERIMENTAL PROCEDURES

Animals

All animal experiments and procedures were conducted in conformity with the Institutional Animal Care and Use Committee (IACUC) guidelines. For liver lipids and mRNA profiling, we analyzed 3-month-old males that were obtained from WT and *Per1/2*^{-/-} mouse colonies derived from the previously described original background (Zheng et al., 2001). Mice were kept under 12 hr light/dark regimen for 2 weeks and fed either ad libitum or exclusively during the dark phase. Throughout the last day, mice were kept under constant darkness and were sacrificed at 4 hr intervals around the clock. Livers were harvested, rinsed in PBS, and rapidly frozen in liquid nitrogen. CT0 corresponded to the time light was turned on and CT12 to the time light was turned off in the animal facility.

Shotgun Lipidomic Analysis

Mice were sacrificed by cervical dislocation; livers were harvested and rinsed with PBS. A ventro lateral section, which corresponds to the right lobe, was cut and instantly frozen in liquid nitrogen. Liver wafers were pulverized into a fine powder by a stainless steel Bio-pulverizer (12 wells, capacity 10–100 mg per well, BioSpec Products) at the temperature of liquid nitrogen. Tissue-fine powders (~15 mg) were weighed from each liver sample and homogenized in PBS by using 2.0 ml cryogenic vials (Corning Life Sciences). Protein levels in the homogenates were quantified using a bicinchoninic acid protein assay kit (Thermo Scientific) with bovine serum albumin as standards. All determined lipid levels were normalized to the protein content of individual samples.

Individual homogenate of the liver samples was accurately transferred into a disposable glass culture test tube. An internal standard mixture for quantitation of all reported lipid classes was added prior to lipid extraction (see Supplemental Experimental Procedures). Lipids were extracted by methyl-tert-butyl ether (Matyash et al., 2008). Each lipid extract was re suspended in a volume of 100 μ l of CHCl₃/MeOH (1:1, v/v) per milligram of protein, flushed with N₂, capped, and stored at -20°C. The whole process of the lipid extraction was performed in a lab with an ambient temperature of 22.7°C \pm 0.2°C and conducted in parallel with the controls.

For ESI direct infusion analysis, lipid extract was further diluted to a final concentration of ~500 fmol/ μ L by CHCl₃/MeOH/isopropanol (1/2/4, v/v/v) with or without 0.02% (v/v) LiOH-saturated MeOH solution, and the mass spectrometric analysis was performed on a QqQ mass spectrometer (Thermo TSQ VANTAGE) equipped with an automated nano spray device (TriVersa NanoMate, Advion Bioscience Ltd.) and operated with Xcalibur software (Han et al., 2008). Identification and quantification of the different lipid molecular species were performed using an automated software program (Yang et al., 2009).

RNA Analysis by Real-Time Quantitative PCR

RNA extraction and transcript quantification by real-time PCR technology were performed as previously described (Asher et al., 2010). Normalization was performed relative to geometrical mean of three housekeeping genes: *Tbp*, *Hprt*, and *Gapdh*. Primers and probes are listed in the [Supplemental Experimental Procedures](#).

Statistics

Data represent mean \pm SEM of four animals per time point. Rhythmicity of lipids was assessed with the nonparametric test, JTK_CYCLE, previously described for the analysis of rhythmic transcripts and metabolites (Hughes et al., 2010). A window of 24 hr was used for the determination of circadian periodicity, and $p < 0.05$ was considered statistically significant. Lipid profiles were crosschecked by visual inspection and false positives excluded.

Measurements of Mice Daily Food Consumption

The daily food consumption of mice was monitored using the Phenomaster metabolic cages (TSE Systems). Measurements of food intake were performed at 15 min resolution throughout the day.

SUPPLEMENTAL INFORMATION

Supplemental Information includes seven figures, six tables, and Supplemental Experimental Procedures and can be found with this article at <http://dx.doi.org/10.1016/j.cmet.2013.12.016>.

ACKNOWLEDGMENTS

We thank U. Albrecht for the *Per1/2* double knockout mice, S. Anpilov for his help with the JTK_CYCLE analysis, and A. Auerbach for his technical assistance with the mouse feeders. The work performed in the laboratory of G.A. was supported by the Israel Science Foundation (ISF), the Abish-Frenkel Foundation, the HFSP Career Development Award (HFSP CDA00014/2012), and the European Research Council (ERC-2011 METACYCLES 310320). The work conducted in the laboratory of X.H. was supported by National Institute on Aging grant R01 AG31675 and by intramural institutional research funds. L.R.-N. and A.N.-C. received a postdoctoral fellowship from the Feinberg Graduate School, Weizmann Institute of Science.

Received: July 27, 2013

Revised: November 12, 2013

Accepted: December 16, 2013

Published: February 4, 2014

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