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Review

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Circadian disruption, clock genes, and metabolic health

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A growing body of research has identified circadian-rhythm disruption as a risk factor for metabolic health. However, the underlying biological basis remains complex, and complete molecular mechanisms are unknown. There is emerging evidence from animal and human research to suggest that the expression of core circadian genes, such as circadian locomotor output cycles kaput gene (*CLOCK*), brain and muscle ARNT-Like 1 gene (*BMAL1*), period (*PER*), and cryptochrome (*CRY*), and the consequent expression of hundreds of circadian output genes are integral to the regulation of cellular metabolism. These circadian mechanisms represent potential pathophysiological pathways linking circadian disruption to adverse metabolic health outcomes, including obesity, metabolic syndrome, and type 2 diabetes. Here, we aim to summarize select evidence from in vivo animal models and compare these results with epidemiologic research findings to advance understanding of existing foundational evidence and potential mechanistic links between circadian disruption and altered clock gene expression contributions to metabolic health-related pathologies. Findings have important implications for the treatment, prevention, and control of metabolic pathologies underlying leading causes of death and disability, including diabetes, cardiovascular disease, and cancer.

Introduction

The daily routines of life on earth are bound to the patterns of the natural environment, leading to the evolution of circadian rhythms in nearly all organisms (1). In animals, circadian rhythms are the daily, timed cycles that coordinate biological functions and prepare the body for recurring diurnal activities such as sleeping, eating, and physical activity (2). Cues such as sunlight, food, sound, and temperature, called zeitgebers, attune circadian rhythms to external conditions, coordinating basic cellular functions to match the environment and maintain homeostasis. A large body of evidence now links an increasing number of known circadian disruptors or zeitgebers, including sleep/wake patterns, dietary timing, and caloric intake, to adverse outcomes in humans. A comprehensive look at evidence from both human and animal studies exploring the relationship between circadian gene expression and metabolic health is lacking. Consequently, we lack a clear mechanistic understanding necessary for future prevention, treatment, and control of metabolic health-related diseases, including diabetes, cancer, and cardiovascular diseases, the leading causes of death and disability across the globe.

Here, we aim to summarize evidence from in vivo animal models and compare these results with epidemiologic research findings to advance understanding of mechanistic links between circadian disruption and altered circadian locomotor output cycles kaput gene (clock) gene expression contributions to metabolic health-related pathologies.

At a cellular level, core clock genes (CCGs) direct the daily oscillatory expression of thousands of clock output genes (COGs) (Figure 1). This complex system regulates a myriad of physiological processes, synchronizing them to match diurnal needs (3, 4). Many of these output genes are associated with other nuclear receptors found throughout metabolic tissues (5). The molecular clock is driven by a dimer of the *CLOCK* and brain and muscle ARNT-Like 1 gene (*BMAL1*) (also known as MOP3 and ARNTL, respectively) transcription factors (6, 7), which binds to genomic response elements known as E-boxes, regulating the expression of thousands of genes, including the period (*PER*) genes (*PER1*, *PER2*, and/or *PER3*) and cryptochrome (*CRY*) genes (*CRY1* and *CRY2*) (8). In turn, the *PER* and *CRY* proteins accumulate in the cytoplasm and, later in the cycle, translocate to the nucleus, where they suppress the *CLOCK:BMAL1* dimer, reducing their own expression (4). Still later in the cycle, *PER* and *CRY* are degraded, *CLOCK* and *BMAL1* expression resumes, and the process repeats with a 24-hour rhythm (9, 10). The *CLOCK:BMAL1* dimer also influences the rhythmicity of the molecular clock through the E-box-dependent regulation of additional transcription factors, such as *RORA*, *RORB*, *RORC*, *NR1D1* (*REVERBA*), and *NR1D2* (*REVERBB*). The *ROR* transcription factors stimulate *BMAL1* transcription, and *REVERB* transcription factors suppress it, creating a second loop that reinforces the E-BOX loop. Together, this transcriptional-translational feedback loop (TTFL) sets off a cascade of transcriptional events that can influence the rhythmic expression of up to 25% of the human genome (2, 4, 9, 11).

In modern society, humans have become less constrained by nature's day/night cycles, and circadian rhythms are pushed and pulled by inconsistent zeitgebers. These circadian disruptors include artificial light, continual food availability, ever-changing work and social demands on sleep/wake timing, and, more recently, diet and

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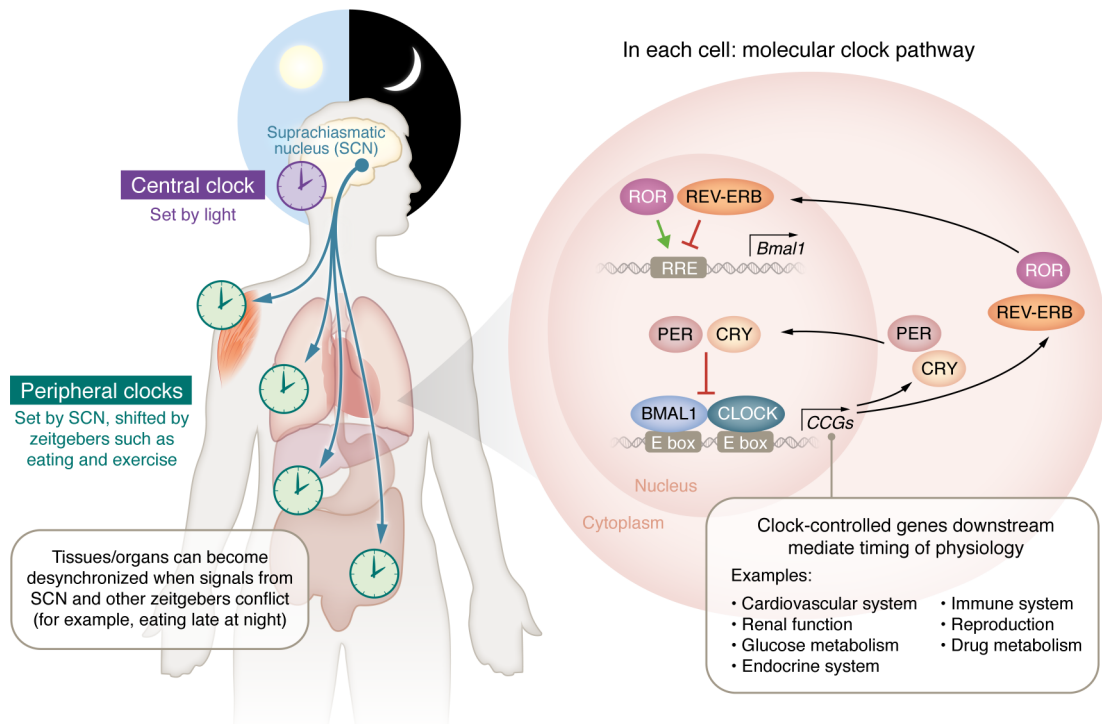


Figure 1. Circadian control of molecular core clock gene signaling and physiologic regulation. The central, peripheral, and molecular clocks and the physiological processes under circadian control. The circadian clock (purple) in the suprachiasmatic nucleus (SCN) of the brain sets peripheral clocks in individual organs and tissue types (light green) via signals including circulating hormones, metabolites, the sympathetic nervous system, and body temperature. Within the cells of the SCN and each organ/tissue type, each cell contains transcription-translation feedback loops, the molecular clocks that drive circadian rhythms. These molecular clocks regulate the transcription of thousands of CCGs and direct the daily oscillatory expression of thousands of CCGs and additional transcription factors that mediate the timing of myriad physiological processes as represented in the molecular clock pathway within cells.

timing of food intake. This disruption is especially pronounced for the approximately 15% of the US workforce who work non-dayshift schedules, at least 35% of whom live in constant sleep deprivation (12, 13). Moreover, studies also report that almost 70% of working adults operate in states of “social jet lag” (14), where circadian disruption occurs as a result of an individual’s shifting or shortening of sleep on weekdays, followed by additional sleep on weekends (15).

When organisms are exposed to circadian disruptors, many biological systems and feedback loops meant to anticipate and regulate diurnal homeostasis can become misaligned through previously described feedback loops at the molecular and cellular levels. Over time, this dysregulation may negatively affect organs and tissues such that catabolic and anabolic processes are out of sync, leading to pathological consequences. For example, aberrant timing of zeitgebers, such as eating during the inactive phase, may induce (a) a generalized environmental desynchrony where the systemic clock is out of phase with external cues such as food timing or (b) an “internal” circadian desynchrony of peripheral clocks (e.g., liver, pancreas) and the central clock, both scenarios with the potential to reduce the efficiency of energy metabolism, leading to weight gain (e.g., see refs. 16–23) and downstream effects on other metabolic pathways. Thus, through regulatory feedback mechanisms, exposure to circadian disruptors may contribute to the development of human disease or inflict adverse health outcomes via inappropriate phase relationships between the internal tissue clock and environmental cues (24–27). Despite relatively rapid rates of behavioral

reentrainment (e.g., sleep patterns) following disruption, internal organs reentrain at different rates, adding additional complexity to the potential impact of circadian desynchrony on human health and disease (28–31). This latter point may be particularly relevant for humans with inconsistent periods of active/inactive cycles over the long term (e.g., shift workers alternating between day and night shifts). Such individuals may rarely reach a state where systemic or tissue clocks are in sync with their external environment, especially energy consumption.

Circadian disruption has been studied as a contributor to the development of the constellation of metabolic health-related pathologies, including obesity, diabetes, and metabolic syndrome (MetS), a cluster of indicators including abdominal obesity, high blood pressure, high blood sugar, high serum triglycerides, and low serum high-density lipoprotein (HDL) cholesterol (32). Early support for this relationship came from both animal models of circadian disruption and human studies of shift workers (33–37). This research suggests that exposure to circadian disruptors leads to mistimed or dampened expression of CCGs and COGs, altering metabolic regulation and contributing to adverse metabolic health and related pathologies (26). Here, we update the current understanding of these phenomena by examining the evidence linking circadian disruption with clock gene expression and metabolism in animal and human studies to provide a foundation for future translational and clinical research applications as summarized in Figure 2 (38, 39).

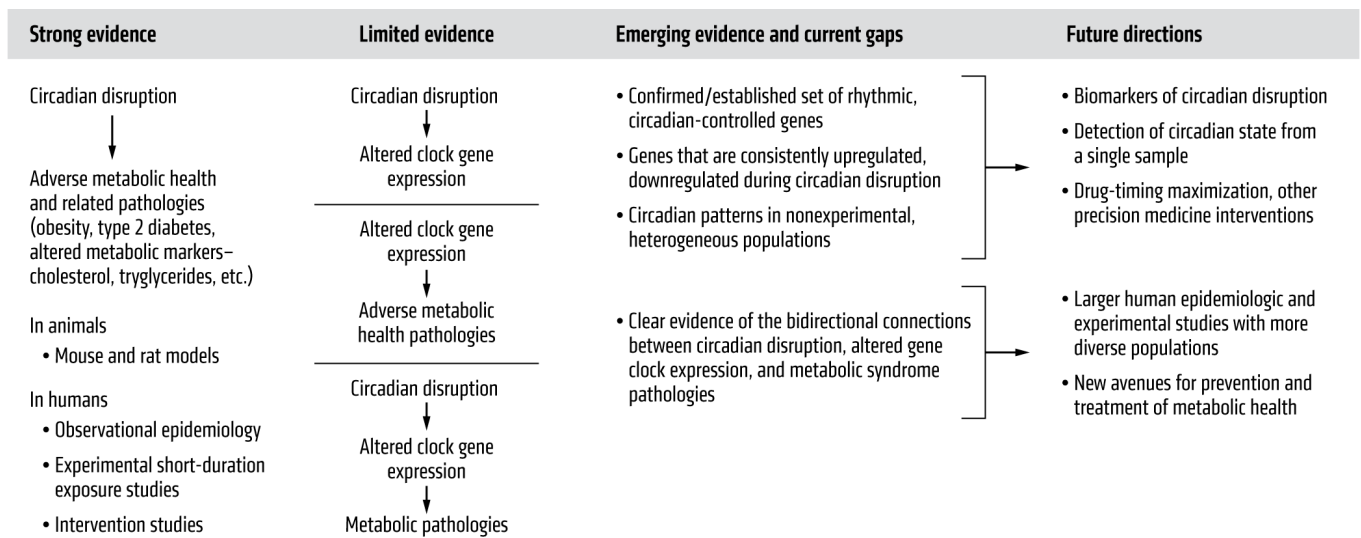


Figure 2. Foundational evidence of core clock gene regulation and metabolic health in animal models and human studies. Summary of the current state of knowledge regarding circadian disruption, CCG expression, and the development of adverse metabolic health–related pathologies.

Circadian disruption, clock genes, and metabolism

In vivo animal models of circadian rhythms. Early investigations into circadian rhythms relied on easily measured physiological aspects of circadian rhythms, such as body temperature and locomotor activity (40, 41). In animals, genetic drivers of circadian rhythms were first revealed through forward genetic screening in *Drosophila melanogaster*, which identified mutants with short, long, and arrhythmic circadian locomotor activity patterns (40, 41). These studies revealed that the *per* gene and its protein product PER, a founding member of the PER-ARNT-SIM (PAS) superfamily, was an essential component of the circadian clock. Subsequent studies in mammals confirmed these *PER* findings and led to the discovery of additional CCGs, *CLOCK*, *BMAL1*, and *CRY1/CRY2*, as well as three orthologs of *Drosophila*: *PER* (*PER1/PER2/PER3*) (37, 38). Notably, *CLOCK*, *BMAL1*, and *PER* were all identified as members of the PAS family of proteins, suggesting hetero- and homodimeric interactions driven by similar domain structures lie at the mechanistic heart of the molecular clock (Figure 1) (5, 36, 37, 42–45).

As mutant models were generated in mice, the circadian clock’s role in metabolic health–related pathologies emerged. Table 1 shows an overview of select studies providing evidence that clock genes are linked to metabolic health pathologies in animal models. For example, in an experiment with mice mutant for the *Clock* gene, mutants displayed altered food intake timing with ad libitum feeding, consuming more calories outside the active phase (37). These animals also had dampened activity rhythms and developed obesity and MetS, including high cholesterol, high triglycerides, high blood sugar, hypoinsulinemia, and elevated leptin during the rest phase. Similar metabolic perturbations were seen in other mouse models with induced mutations in molecular clock components. For example, *Bmal1*-null mice displayed blunted postprandial insulin responses, decreased gluconeogenesis, and loss of typical glucose and triglyceride rhythms (36, 45). Likewise, *Per2*-mutant mice developed without normal glucocorticoid rhythms (46) and *Cry1/Cry2* double mutants displayed altered liver metabolism and altered patterns of circu-

lating growth hormone (44). Finally, when compared with WT, transgenic mice generated to overexpress *Cry1* exhibited greater hyperglycemia without increased weight gain (47).

Further examinations designed for understanding the relationship between clock and glucose homeostasis in *Clock*-mutant mice have revealed altered expression of genes involved in pancreatic islet cell development and insulin signaling (45). At eight months, these mutants displayed elevated serum glucose compared with WT counterparts, seemingly consequent to defective glucose-stimulated insulin release from the *Clock*-mutant islets. Interestingly, findings from young *Clock*-mutant mice aged two to three months demonstrate that a compensatory component exists early on, driven by the *Clock* mutation in other organ systems such as the liver, leading to age-related emergence of insulin resistance and underlying deficits in insulin secretion manifested as pathogenic hyperglycemia (i.e., diabetes) (45). An explanation for the age-dependent hyperglycemia phenotype may come from the observation that *Clock*-mutant and *Bmal1*-null mice exhibit loss of typical glucose and triglyceride rhythms, with impaired and abolished gluconeogenesis, respectively (36). Because gluconeogenesis occurs predominantly in the liver, the contrasting effects between the paired metabolic organs regulating glucose homeostasis in *Clock*-mutant and *Bmal1*-null mice caused a “masking” of the phenotype when the animals were young. Examples such as this highlight the challenges in disentangling the role of the circadian clock when using global mutant/null models (36).

To address this issue, pancreas-specific *Bmal1*-null mice (*Bmal1^{fl/fl}; Pdx1-Cre*) were employed to knock out the circadian clock in pancreatic islet cells. These experiments revealed that, even when mice are at a young age, the islet cell clock influences insulin secretion, glucose levels, and glucose tolerance (i.e., hypoinsulinemic diabetes) (45). Thus, the discrepancy in young versus old *Clock*-mutant mice may be due to the circadian clock yielding different tissue-specific effects in metabolic processes such as glucose metabolism, which is particularly relevant for the pathologic processes of diabetes and MetS. Further, many metabolic condi-

Table 1. Summary of evidence of support for the role of circadian disruption and CCGs in regulating metabolic health from animal studies

CCGs studied	Tissue/animal model	Outcomes/mechanisms examined	Conclusions and advancements	References
Genetic evidence from knockout and transgenic mouse experiments for gene expression control of adverse metabolic health outcomes				
Clock	Mutant mouse with altered clock gene expression	Altered diet Cholesterol, high triglycerides, high blood sugar, and hypoinsulinemia, and elevated leptin during rest phase	<i>Clock</i> genes control diet (27) and activity and, in parallel, key metabolic indicators	(37)
Bmal1	<i>Bmal1</i> -null mice	Altered insulin response, gluconeogenesis	<i>Bmal1</i> controls metabolic pathways, including glucose and insulin regulation	(36, 45)
	<i>Per2</i> mutant		Disrupted glucocorticoid rhythms	(5, 97)
Cry1/Cry2	<i>Cry1/Cry2</i> mutants	Liver metabolism Circulating growth hormone	<i>Cry1/Cry2</i> mutants influence liver metabolism and hormone homeostasis	(44)
	Transgenic <i>Cry1</i> overexpression		<i>Cry1</i> is important in glucose regulation, including in diabetes phenotypes	(47)
Clock gene expression, compensatory organs, and glucose metabolism				
Clock	Mutant mouse	Compensatory mechanisms in young compared with old mice showed role of other organs including liver	Insulin resistance in old mice (>8 months) emerges over time	(45)
Bmal1	Young <i>CLOCK</i> -mutant and <i>Bmal1</i> -null mice compared with older mice	Altered liver and pancreas leads to impaired gluconeogenesis	Role of global mutant and null models shows interdependence in tissues and need for further examination	(36)
Bmal1	Pancreas-specific <i>Bmal1</i> -mutant mice (<i>Bmal1^{fl/fl}; Pdx1-Cre</i>) to knock out the circadian clock in pancreatic islet cells	Insulin secretion in relation to altered clock gene expression in pancreatic islet cells and longitudinal changes with age in mutant mice	Further confirms that circadian clock impacts on glucose metabolism, particularly disruption of the pancreatic clock, contribute to diabetes via alterations of insulin secretion that can change with age	(45)
Bmal1	Pancreas-specific <i>Bmal1</i> -mutant mice (<i>Bmal1^{fl/fl}; Pdx1-Cre</i>) to knock out the circadian clock in pancreatic islet cells	A role for pancreatic islet cell clock-driven rhythmic expression controlling insulin secretion	Further evidence of a tissue-specific response governed by core sets of genes that regulate rhythms in all tissues	(52)
NRs (related to CCGs)	C57BL/6J mice	Differential and rhythmic expression of numerous nuclear receptors in metabolic tissues	Further evidence that a circuit of NRs is regulated across multiple tissues related to CCGs and metabolic health and is associated with lipid and glucose metabolism	(5)
External exposure to circadian disruptors (zeitgebers: light, diet, timing of food intake) and impact on metabolic health				
Phase shift	WT Sprague-Dawley and diabetes-prone HIP rats	WT rats exposed to a phase-shift or constant-light protocol did not experience changes in metabolic markers, including blood glucose control, β cell function, mass, turnover, and insulin sensitivity; however, rats with reduced β cell functioning did	β Cell functioning plays an important role in altering influence of light disruption among rats and when deficient can accelerate development of T2D	(58)
	Male and female WT C57BL/6J (WT) mice	A phase-shifted chronic jet lag protocol led to altered rhythmicity of CCGs. Alterations in downstream genes involved in insulin regulation were also observed	Approximately 5% of the transcriptome in the pancreas of mice is regulated by CCGs, and external phase shifting alters regular rhythmic control of genes in the pancreas associated with insulin and enzyme regulation	(56)
TRF	Mice fed high-fat diets and under TRF protocol	Protective effects of TRF on numerous MetS outcomes, including impaired glucose tolerance, insulin sensitivity, and inflammation.	High-fat diets change both clock gene expression and amplitude, but can be restored via TRF even on high-fat diet, leading to protective phenotypes	(59-61)
Timing of food intake	Mice fed high-fat diets and under TRF protocol	Thermogenesis in adipocytes with respect to negative metabolic consequences: weight gain, glucose tolerance, insulin sensitivity, and inflammation	CCGs regulated mediators responsible for establishing controlled thermogenesis in adipocytes, the primary mechanism underlying MetS pathologies associated with diet and clock genes	(59)
Phase shift and TRF	Mice, light, and food	COG and SCN synchrony and metabolic health (weight gain, hormone changes) in relationship to exposure to light and food in mice	While light is a primary circadian disruptor, food intake and timing can also support synchronization and desynchronization, and both light and food intake influence metabolic outcomes in an interdependent manner	(56-58)

NR, nuclear receptors; HIP, human islet amyloid polypeptide transgenic.

tions emerge in humans with aging, e.g., type 2 diabetes (T2D), nonalcoholic steatohepatitis, etc. Concordantly, disruption of the circadian clock in humans results from global disruption/desynchrony. Therefore, it is plausible that the early impact of circadian desynchrony on MetS in humans is difficult to completely understand given the organ-specific functions of the clock. In fact, disease states may only become apparent after long-standing desynchrony, which is supported by studies demonstrating increased risk of developing diseases such as diabetes with increasing duration of shift work (48–51).

Glucose homeostasis is influenced by cellular-specific gene signaling mechanisms controlled by specific CCGs. For example, a 2015 study by Perelis et al. examined pancreatic β cells from mice with intact or disrupted BMAL1 expression. For intact cells, CLOCK/BMAL1 dimers were shown to bind to regulatory sites (CCGs) in islet cells to drive transcription of genomic targets (COGs) in these cells that were distinct from those of other cells in the liver. They further found that mice with disrupted BMAL1 expression developed glucose intolerance, suggesting a direct genetic mechanism controlling diabetes as one measure of metabolic health (52). Further, the majority of CLOCK/BMAL1 binding sites identified within β cells are not commonly identified in the liver or other tissues — supporting the tissue-specific role of the clock in altering metabolic health through active enhancer regions and epigenetic chromatin regulation of unique genes within cells and suggesting that polymorphisms or alterations contribute to metabolic disruption. Tissue-specific subsets of circadian genes reinforce this idea. Despite the substantial number of cycling genes in each tissue (5), only a small number of common genes are rhythmic in all tissues (11).

A large body of work has been undertaken for understanding the consequences of external/environmental circadian disruptors in mice (53). These protocols are meant to mimic circadian disruptors commonly experienced by humans, such as altered timing of light exposure, activity, sleep, or food intake (17, 30, 54–56). Experimental studies aiming to mimic shift-work exposures through phase-shift and time-restricted feeding protocols suggest that exposure to these various circadian disruptors can alter metabolic health and CCG expression (56–58). A 2011 study found that phase-shift protocols among Sprague-Dawley rats were associated with changes in the acceleration of multiple indicators of T2D, particularly for animals with altered β cells (58). Similarly, a 2021 study of circadian disruption via chronic jet lag investigated transcriptional changes in mice and found that 5% of the transcriptome in the pancreas is regulated by CCGs and that external phase shifting in mice alters regular rhythmic control of genes in the pancreas associated with insulin and enzyme regulation (56).

Emerging evidence from experimental food-intake and timing models illustrates the multiple mechanisms by which circadian disruption affects metabolic health. For example, WT mice subjected to misaligned food intake relative to active and inactive periods display accelerated weight gain similar to that of humans. Along with lowered amplitude of clock genes, mice consuming a high-fat diet during their inactive period gained substantially greater weight and had a higher body fat percentage than mice consuming the same diet during their active period (18). In contrast, when a time-restricted feeding (TRF) protocol restricted feeding to the active period, mice on high-fat diets were protect-

ed from weight gain and the increased markers of adverse metabolic health experienced with ad libitum feeding. In several studies, TRF prevented obesity and impaired glucose tolerance, restored insulin sensitivity, and protected against inflammation and hepatic steatosis (59–61). Moreover, during the active phase, TRF restored normal hepatic glucose metabolism elicited by the ad libitum high-fat diet. TRF also restored the oscillation of metabolic regulators in the liver that were dysregulated with an ad libitum high-fat diet. Moreover, it restored CCG expression amplitude. Thus, eating at times misaligned to circadian rhythms leads to altered metabolic health, while TRF ameliorates this pathology.

A combination of TRF and diet quality affects metabolic health by regulating CCGs, as seen in a series of critical mouse experiments. Animals were subjected to high-fat diets with either ad libitum feeding, TRF confined to an active period, or TRF confined to an inactive period (59). Despite similar activity and timing of food intake overall, a high-fat diet during the inactive period caused increased body mass and lower energy expenditure. Carbon-labeling studies demonstrated that the high-fat diet during the inactive period resulted in decreased glycolysis in adipocytes and dampened oscillation of CCGs compared with the high-fat diet during the active period. A series of genetically engineered mouse models demonstrated that the adverse metabolic consequences of a high-fat diet during the inactive period were related to impaired adipocyte thermogenesis. The mediators of thermogenesis in the adipocytes were found to be regulated by the core molecular clock and responsible for maintaining metabolic health in the high-fat diet/active-phase TRF group. This study supports the idea that optimal metabolic health can depend on the alignment of food intake and the biological rhythms of cellular thermogenesis controlled by the molecular clock. Moreover, these data help us to understand the contributions of high-risk dietary regimens and circadian misalignment. These results also underscore important considerations for humans experiencing circadian disruption, leading to the suggestion that mitigation of risk for adverse metabolic health could include strategies to align specific nutrient intake with the internal clock and highlighting the importance of developing simple and reproducible methods of monitoring organ-specific reentrainment that may facilitate optimizing such an approach.

Foundational epidemiologic and population-based research findings. While human studies often lack the specificity to advance mechanistic insights, coupled with animal-based findings, they provide crucial foundational evidence linking circadian disruption to metabolic health, as summarized in Table 2. Some of the first studies to suggest circadian disruption alters metabolic health come from epidemiological investigations of shift workers. For example, an examination of 54,724 participants in the Nurses' Health Study II found that individuals exposed to any duration of night-shift work had increased odds of obesity, higher total calorie intake, and shorter sleep durations than those who had never worked night shifts, after adjusting for age and socioeconomic status (62). In a separate study, shift workers displayed increased odds of being overweight or obese and were more likely to report insufficient sleep than individuals working traditional schedules. Moreover, there was a stronger association between shift work and overweight conditions among individuals reporting insufficient sleep, suggesting a protective effect of adequate sleep dura-

Table 2. Human exposure to circadian disruptors (zeitgebers: light, diet, timing of food intake) and influence on adverse metabolic health outcomes: results from select epidemiologic investigations and experimental studies

Exposure	Key sample characteristics	Metabolic health outcome(s)	Other related outcomes	Conclusions	References
Shift work	54,724 Participants in the Nurses' Health Study II	Increased odds of obesity	Higher calorie intake, shorter sleep durations	Results showed differences in metabolic health and metabolic health-related behaviors after adjustment of income and age effects	(62)
	Representative statewide sample of 1,593 Wisconsin adults	Increased odds of being overweight	Increased odds of sleep disturbance	Influence of shift work on sleep patterns and obesity	(63)
	Twenty-four overweight women, divided into night-shift workers (<i>n</i> = 12) and day-shift workers (<i>n</i> = 12)	Greater fat mass, lower insulin sensitivity, higher triglycerides, altered dietary enzymes	Increased energy intake, impaired sleep are also associated with night shift	Shift work may directly influence metabolic health and indirectly be associated through altered appetite regulation; however, study sample is small	(64)
SJL, chronobiology, and dysregulated sleep	815 individuals from the Dunedin longitudinal study	SJL associated with metabolic indicators, including elevated BMI, fat mass, and MetS and increased waist circumference	All analyses controlled for several lifestyle factors; SJL was also associated with elevated HbA1c in metabolically unhealthy obese individuals	In observational studies of humans, circadian disruption via SJL is associated with metabolically unhealthy phenotypes	(65)
	490 Midlife men and women from AHAB-II	Multiple indicators of metabolic health were adversely associated with SJL, including lower HDL cholesterol and higher triglycerides	Chronotype alone associated with lower HDL cholesterol	Many markers of metabolic health were altered among individuals with objectively measured SJL independently of chronotype or propensity for morning or evening activity, suggesting circadian disruptors, including light and sleep regularity, may contribute to misalignment linked with metabolic health	(66, 67)
	SWAN sleep study of 338 women	Greater bedtime variability and greater bedtime delay associated with altered insulin resistance (homeostatic model assessment-insulin resistance) and BMI	Results were partially explained by SJL or later bedtimes on weekends		(68)
	191 Community-dwelling adults over age 80 years	Objectively measured variations in sleep timing and patterns were associated with increased obesity and altered body composition	Results remained consistent after adjustments for numerous markers of sleep variability simultaneously	At older ages, sleep variability measured in numerous ways, as indicators of potential circadian misalignment, are also associated with indicators of adverse metabolic health	(69)
Experimental SJL protocols	21 Healthy adults (10 men and 11 women) imposed 28-hour days and 5.6 hours per night sleep restriction	Increased fasting and postprandial glucose levels and significantly decreased resting metabolic rates, reduced insulin sensitivity, and weight gain	In one, changes normalized after 9 days of return to normal sleep/wake patterns; in another experiment, dietary patterns also changed	Experimental studies showed that short-duration sleep disturbances (10 days) also lead to adverse changes in metabolic indicators; however, a return to regular patterns may reverse impacts. Sleep disturbance in one study also contributed to changes in dietary habits, making it difficult to disentangle circadian disruption from other behavioral factors	(70)
	Three groups of healthy young adults were sleep restricted for two 5-day weeks of 5 hours per night, 1 weekend catch-up group, 1 control	Reduced insulin sensitivity and weight gain among sleep-restricted groups	Individuals with disrupted sleep protocols also increased nighttime food intake	Sleep disturbance in one study also contributed to changes in dietary habits, making it difficult to disentangle circadian disruption from other behavioral factors, and weekend catch-up did not rectify minimized short-duration sleep	(71)
TRF	Numerous TRF clinical trial studies (19) reviewed in meta-analysis	TRF reduced body weight, decreased fat mass, preserved fat-free mass, reduced systolic blood pressure, triglycerides, and fasting glucose		In experimental studies, TRF may reduce adverse metabolic health outcomes and may be considered a successful mechanism to counteract circadian misalignment	(72)
Sleep and timing of food consumption	19 Healthy younger adults including 12 men and 7 women with mean age 26; 14-day sleep restriction and restricted daytime eating	Endogenous circadian glucose rhythms associated with circadian disrupted sleep and modified by meal timing		Sleep and timing of food consumption may be very interrelated in driving circadian misalignment and metabolic health	(73)

SJL, social jet lag; AHAB-II, Adult Health Behavior Project Phase-2; SWAN, Study of Women's Health Across the Nation.

tion during shift work (63). A much smaller study of 24 women (12 night-shift and 12 day-shift workers) found that night-shift workers had greater fat mass, larger energy intake, impaired sleep, lower insulin sensitivity, and higher triglycerides compared with their day-worker counterparts (64). Night-shift workers also had higher postprandial ghrelin levels and lower bloodstream levels of xenin, a gut-derived hormone, offering preliminary evidence of shift work and metabolic health in the form of appetite regulation (64).

Another significant circadian disruptor linked to metabolic health is “social jet lag,” measured as the difference in midsleep time on nights before work or school and those before work- or school-free days (14). One study of 815 nonshift workers born between 1972 and 1973 in New Zealand identified an association between social jet lag and numerous metabolic health indicators, including BMI, fat mass, and waist circumference (65). In independent studies, individuals with social jet lag displayed greater adiposity, lower HDL cholesterol levels, higher triglycerides, increased insulin resistance, and higher fasting plasma insulin even after controlling for behavior and sleep quality. Moreover, individuals with a tendency to be most active in the evenings and delay sleep onset, known as “evening chronotypes,” had lower HDL cholesterol levels, consistent with similar findings that evening types have a heightened risk for cardiovascular disease, given their proclivity to circadian-disrupted schedules and social jet lag in particular (66, 67).

Additional insights come from several studies among non-shift-work female populations. A cross-sectional analysis of non-shift-working middle-aged women revealed a positive association between bedtime variability and bedtime delay with increased insulin resistance (68). After more than five years of follow-up, greater bedtime delay predicted higher insulin resistance, suggesting that both acute- and chronic-inconsistent sleep timing induce metabolic dysfunction (68). These observations were echoed among a study of older women over the age of 80 that found intraindividual variation in objectively measured wake time, sleep duration, sleep delays (social jet lag), and the midpoint of sleep were associated with alterations in body composition, including percentages of increased fat mass and lower lean mass as metabolic health indicators (69).

Experimental phase-shift studies among adults requiring short-term alterations of regular sleep patterns provide additional evidence for links between circadian disruptors and markers of metabolic health. In one such study, 21 healthy adults (10 men and 11 women) of varying ages were subjected to circadian disruption, achieved via imposed 28-hour days and 5.6 hours per night sleep restriction. After three weeks, circadian-disrupted participants had increased fasting and postprandial glucose levels and significantly decreased resting metabolic rates compared with baseline, a pattern observed in both young and old subjects. These changes normalized after nine days of return to standard sleep/wake patterns (70). In a separate study, two experimental groups of healthy young adults were exposed to two five-day weeks of five hours per night of sleep restriction, with one group permitted an interim two-day weekend of ad libitum “catch-up” sleep, while the other group continued sleep restriction for the entire study period. A control group with normal sleep (nine-hour daily sleep windows) was also included. Despite sleeping an extra three hours each weekend day, the weekend catch-up group experienced reduced insulin sensitivity relative to controls.

While all three groups increased their energy intake compared with baseline calorie-controlled meals, only the control group did not experience weight gain. Moreover, both disrupted groups increased after-dinner snacks on days following sleep restriction. These findings suggest that a weekend catch-up sleep after sleep debt, akin to social jet lag, is not sufficient to correct the metabolic dysfunction accrued during sleep restriction (71).

There is mounting evidence that TRF, particularly TRF during the active phase and aligned with circadian rhythms, is also beneficial for human metabolic health, which is similar to observations in mice. A meta-analysis of 19 TRF clinical trials found TRF diets significantly reduced body weight, decreased fat mass, preserved fat-free mass, and reduced systolic blood pressure, triglycerides, and fasting glucose (72). TRF is thus seen as a promising treatment for changing metabolic health, even without reducing the total calorie intake, hence emphasizing the importance of circadian timing in metabolic processes (72). Further, in a disrupted sleep and TRF study, participants with three days of 28-hour day simulated night-shift protocols were divided into two groups and fed meals in sync with their shifted/disrupted schedule or their usual dietary intake schedule. The group fed in sync with their shifted/disrupted, 28-hour cycle (eating meals between midnight and 4 am, when they would typically be asleep) showed impaired glucose tolerance (73). In parallel with experimental animal findings, human study results further establish the role of food intake as a strong entraining cue for the circadian clocks in the periphery and indicate metabolic outcomes can be improved by aligning the food-intake window with the active phase while removing windows of food intake later in the day or night.

Circadian disruption and CCG expression. Circadian disruption is also associated with altered clock gene expression in human studies, offering clues to the biological mechanisms linking circadian disruption with metabolic health. A small 2019 study including 18 female nurses revealed fewer rhythmic genes observed in the PBMCs of rotating night shift workers ($n = 9$) compared with nurses working day shift ($n = 9$). Moreover, phase desynchrony of core body temperature, peak cortisol, and dim-light melatonin onset were also observed (74). A more extensive investigation of 60 nurses using a single measurement time point identified differential expression of nearly all CCGs, including transcripts of the CCGs *BMAL1*, *CLOCK*, *NPAS2*, *PER1*, *PER2*, *PER3*, *REVERBA*, *CRY1*, and *CRY2* genes. These CCGs were dysregulated in morning blood draws taken among rotating shift nurses and compared with those day-shift nurses (75). A small study of eight participants exposed to a simulated night-shift work protocol, including three days of ten-hour sleep period delay, repeated blood draws over two 24-hour periods, and transcriptome-wide analysis of PBMCs, showed a significant reduction of rhythmic transcripts, including *PER3* postdisruption compared with baseline (76).

Interestingly, similarly to what was found in the 2019 nurses' study (74), some CCGs, including *PER1* and *BMAL1*, maintained rhythmicity. Genes that remained rhythmic after disruption displayed dampened oscillation patterns that matched habitual sleep and wake times rather than the newly disrupted schedule. These results show that, while circadian function may remain largely intact in a simulated night shift, many COGs lost expression amplitude, influencing several important metabolic pathways, including immune-system regulation (76).

Table 3. Transcriptome-wide changes and circadian clock gene expression linked among select human studies

Exposure	Key sample characteristics and study design	CCG expression	Metabolic health-related outcomes	Conclusions	Reference
Shift work	18 women+, 9 shift workers compared with 9 nonshift workers: 3-hour blood draws, PBMC analyses	Transcriptome-wide PBMC analysis showed alterations in circadian transcription and phase shifting of CCGs between day- and night-shift workers, with the exception of <i>PER1</i>	Phase desynchrony of core body temperature, peak cortisol, and dim-light melatonin onset was also observed	Transcriptional changes in oscillation of clock genes comparing a small sample of day- and night-shift workers was consistent with previous investigations of circadian misalignment. Interpretation of results is limited due to small sample size	(74)
	60 shift workers, single morning blood draw, comparing rotating day and night shift workers	Differential expression of nearly all CCGs	Transcriptome-wide changes	Women who are regular night-shift workers compared with women who regularly work during the day had differentially expressed CCGs during morning draws. It was unclear whether this is due to blood draw time of day or influence of prolonged shift work	(75)
	8 participants+, 10-hour sleep delay, 24-hour repeated blood draw, PBMC analyses	Significant reduction of rhythmic transcripts including <i>PER3</i> after disruption, and many CCGs lost expression amplitude; notably <i>PER1</i> and <i>BMAL1</i> maintained rhythmicity	Genes in metabolic pathways including immune-system regulation also dysregulated from baseline	Transcriptome-wide analyses showed rapid shifts in oscillation and rhythmicity of CCGs, with some exceptions (<i>PER1</i> and <i>BMAL1</i>); nonetheless, other gene transcripts affecting metabolic health were also dysregulated, suggesting circadian misalignment alters gene expression beyond just CCGs	(76)
Sleep-induced circadian disruption	22 healthy adults, sleep-restricted protocol shifting sleep by 4 hours for 3 days	Loss of rhythmicity across the transcriptome, including in CCGs <i>CLOCK</i> , <i>ARNTL</i> , and <i>PER3</i>	Differential expression in metabolic pathways among those with forced desynchrony	Forced sleep desynchrony	(77)
Sleep deprivation	26 healthy adults, exposed to 1 week of insufficient sleep (5.7 hours) compared with controls (8.5 hours of sleep)	Differential expression of CCGs observed including <i>PER1</i> , <i>PER2</i> , <i>PER3</i> , <i>CRY2</i> , <i>CLOCK</i> , <i>NR1D1</i> , <i>NR1D2</i> , <i>RORA</i> , <i>DEC1</i> , <i>CSNK1E</i>	Differential expression in metabolic pathways among those with insufficient sleep	Sleep deficiency also influenced CCG expression, and key metabolic-related processes across the entire transcriptome	(78)

In a separate forced desynchrony experiment, 22 participants underwent baseline assessment followed by several 28-hour days of forced desynchrony, with sleep onset pushed back four hours further each night. During disruption, 24-hour melatonin rhythms remained largely preserved as compared with baseline rhythms. In contrast, there was a six-fold reduction in PBMC-measured circadian transcripts, including *BMAL1*, *CLOCK*, and *PER3* (77). Experimental sleep-deprivation studies also aimed at identifying resultant transcriptome alterations have identified similar reductions in the circadian rhythmicity of genes and changes in the expression of genes from chromatin-remodeling immune and stress-response pathways. In a study of 26 sleep-restricted individuals who were limited to six hours per night, gene expression in thousands of transcripts was altered, and the number of genes with detectable circadian patterns was reduced by 20%. Again, *CLOCK*, *PER1*, *PER2*, *PER3*, *CRY2*, and *RORA* were significantly impacted. Additionally, several metabolic and oxidative stress-related genes were altered after the sleep-restriction protocol (78). It is important to note that these transcriptome-wide analyses were conducted on relatively small numbers of individuals with short desynchrony and sleep-deprivation protocols. Therefore, the longer-term impacts of these exposures are unclear. Nonetheless, these findings collectively support the idea that circadian misalignment has widespread effects on the transcriptome and has a differential impact on central versus peripheral clocks, highlighting the potential for adverse metabolic health from disruption of the habitual sleep/wake cycle (see a summary of findings in Table 3).

CCGs and metabolic pathology. The connection between clock gene expression and metabolic health may be bidirectional. Evidence from various human studies suggests that a state of adverse metabolic health itself, including elevated BMI, can alter the expression of CCGs in a tissue-specific fashion. In one study of 21 lean and 28 morbidly obese female nonshift workers without diabetes, examination of 24-hour adipose gene expression revealed that obese individuals displayed altered circadian expression of many CCGs, including *CLOCK*, *BMAL1*, *PER1*, *CRY2*, and *REVERBA*, compared with healthy, lean subjects. Positive correlations were found among all subjects between *REVERBA* and BMI/waist circumference, *CLOCK* and LDL cholesterol, and *RORA* with HDL cholesterol. An interesting conclusion from this study was that *REVERBA* is an important gene associated with metabolic health (79).

Further evidence of connections between metabolic phenotypes driving circadian disruption comes from studies showing that weight loss alters CCG expression patterns. A 2020 study examined differential mRNA levels and expression of CCGs in skeletal muscle among 23 obese patients (5 women and 18 men) undergoing gastric bypass surgery and 14 normal-weight controls (6 women and 8 men). Males in the obese group had significantly lower *CLOCK*, *CRY1*, and *CRY2* expression than lean male controls (80). Obese women exhibited downregulated *CRY1* mRNA levels compared with lean female controls, but *CRY1* expression was restored to lean-control levels following gastric bypass-induced weight loss (74). Interestingly, while changes in CCG expression varied by sex, additional research is needed to replicate these findings. A second study examined participants'

expression of CCGs in adipose tissue before and after hypocaloric diet-induced weight loss (81). After eight weeks, 50 subjects who lost 8% or more of their body weight saw significant increases in *PER2* expression compared with baseline, with similar changes to genes regulating fat metabolism, autophagy, and inflammatory responses (81).

Human studies also suggest that even limited, short-term exposure to circadian disruptors can alter metabolic pathways and clock gene expression. One such study, in 2018, subjected 14 healthy men to three days of normal sleep, followed by three days of reversed day/night schedules. After three days of disruption, fasting glucose and free fatty acids were significantly elevated compared with what occurred with normal sleep conditions (82). In addition, a significant transcriptional alteration in PPAR signaling was observed, leading to the hypothesis that misalignment promotes a preference for intramuscular fatty acid metabolism over glucose metabolism. Interestingly, after the three days of misalignment, CCGs had not reentrained to the reversed schedule and remained aligned to the regular day/night schedule (82). In another 2015 investigation, 15 healthy male participants were exposed to acute 24-hour sleep deprivation, and increased methylation in the *CRY1* and *PER1* genes in adipose tissue was observed, suggesting that methylation is also a mechanism for the downregulation of CCG expression (83). Furthermore, after sleep deprivation, expression of *BMAL1* and *CRY1* in skeletal muscle was decreased, and postprandial plasma glucose concentrations were increased (83). These studies also provide evidence that even short-term misalignment of the circadian clock from standard behavior patterns, from a single night of wakefulness to a few days of misalignment, can be linked to metabolic changes in humans.

Studies of CCG expression have also shown that TRF studies can help combat circadian misalignment's negative consequences, suggesting new opportunities for preventing and treating adverse metabolic health outcomes. In a 2019 crossover study of 11 obese participants, the efficacy of TRF was investigated (84). Comparison of four-day ad libitum feeding (8 am–8 pm) with an early daytime feeding window (8 am–2 pm) revealed that TRF resulted not only in increased expression of CCGs (*BMAL1*, *CRY1*, *CRY2*, and *RORA*), but also elevated ketones, elevated cholesterol levels, reduced mean blood glucose levels, and reduced glucose spikes throughout the day. This was despite equal calories consumed between conditions. While many studies have supported using TRF to improve metabolic health, this experiment provided key insights by measuring and associating CCG expression with improvements in lipid metabolism and glucose regulation (84). A similar randomized case-crossover study of TRF examined 11 men who were either overweight or obese and found that TRF improved daytime insulin profiles and reduced night-time glucose levels. The oscillation patterns of CCGs, including *CLOCK*, *BMAL1*, *CRY1*, *PER1*, -2, and -3, and *REVERBA* and -B, were unchanged between 15-hour free-feeding and 8-hour TRF conditions (85). However, the authors identified an increase in the amplitude of oscillating muscle transcripts related to amino acid transport, suggesting that TRF has multiple health benefits (85).

As in mice, genetic variation in CCGs and genes that modify the clock can influence rhythms and metabolic health in humans. For example, in a meta-analysis of cohort studies from the Cohorts for Heart and Aging Research in Genomic Epidemiology Consor-

tium, associations among sleep duration, genetic variants of core clock and other circadian genes, and cardiometabolic traits were identified (86). In carriers of the T allele of the melatonin receptor 1B, long sleep duration (9+ hours per night) was associated with increased BMI. Additionally, in carriers of the A allele of *CRY2* (SNP rs11605924), sleep duration was positively associated with HDL cholesterol level (86). Additional evidence supporting the connection can be seen in investigations of SNP mutations and their role in metabolic health, reviewed in Škrlec et al. (87). The growing evidence of CCG SNPs underlines the complex relationships between CCGs and metabolic health. Table 4 summarizes several examples and study findings.

Conclusions

Research across organisms supports the idea that CCGs and clock-controlled genes control link circadian disruption and metabolic health, including MetS and diabetes. Specifically, there is a substantial and growing body of evidence from CCG mutant mouse models linking clock activities to pathological metabolic outcomes. Tissue-specific CCG mutant mouse models have also helped elucidate the effects of CCGs in each organ and tissue type. Moreover, TRF protocols in mice offer strong evidence for food intake as a zeitgeber/disruptor and consequential CCG changes that have metabolic pathological consequences.

From a risk paradigm, circadian disrupter impacts are a consequence of their quality and exposure duration. Alternating shift work (as defined by discrete and alternating periods of daytime, swing, and night work) provides an example of this concept. In a tissue desynchrony model, the oscillating metabolic profiles of distinct tissues are out of proper alignment due to the differential response times required to reestablish normal rhythms after an inappropriate zeitgeber exposure or significant circadian perturbation. Extended periods of alternating shifts may not allow for appropriate internal organ reentrainment. This has potential relevance in alternating shift workers and other highly dyssynchronous populations, such as frequent international travelers or with weekly social jet lag. The continual shift in zeitgebers likely requires a unique strategy to optimize realignment and promote a positive impact on health, including mitigation of adverse metabolic pathologies.

It may also be important to emphasize that shift work is not always synonymous with circadian desynchrony. One can predict the existence of many shift workers with appropriate circadian hygiene that allows alignment of the timing of nutrient intake with internal physiologic processes (88). The challenge lies in how to disentangle those components of circadian misalignment contributing to poor human health, given the potential for synergy, additivity, or antagonism for a spectrum of prevalent disruptors such as extended timing of food intake (i.e., eating outside the active phase) and extended exposure to light at night (screen time) intertwined with poor sleep habits.

With these ideas in hand, it becomes possible to more precisely define circadian disruptors as those risk factors that lead to circadian desynchrony. In a simple example of this idea, circadian disruption results from the inappropriate timing of a zeitgeber, which can alter the clock phase, decrease its amplitude, and disrupt synchrony in core clock factors and outputs

Table 4. Genetic polymorphism circadian clock gene(s) linked to metabolic health in select human epidemiologic and experimental studies

CCGs studied	Population characteristics	Metabolic health outcome(s)	Gene-environment interactions examined	Conclusions	Reference
BMAL1 PER3 RORA RORB	3000 Taiwanese adults	MetS, as defined by the International Diabetes Foundation, comprises altered values on a cluster of metabolic indicators, including insulin resistance, raised blood pressure, abdominal obesity, and dyslipidemia (raised triglycerides, lowered HDL cholesterol)	Alcohol consumption, physical activity, smoking	No significant associations between core clock SNPs and MetS after Bonferroni's correction Significant interactions between SNPs in <i>BMAL1</i> , <i>GSK3B</i> , <i>PER3</i> , <i>RORA</i> , <i>RORB</i> on MetS Interactions between SNPs in <i>BMAL1</i> , <i>PER3</i> , <i>RORB</i> , all measured health behaviors, and MetS incidence	(98)
CLOCK	7,098 adults aged 55–80 with 3+ cardiovascular risk factors or T2D	T2D Cardiovascular disease events (stroke, myocardial infarction, cardiovascular death)	Mediterranean diet	G allele of <i>CLOCK</i> -rs4580704 SNP had lower incidence of T2D compared with CC homozygotes; Mediterranean diet increased protection for G allele carriers In T2D subjects, G allele carriers have increased stroke risk vs. CC homozygotes	(99)
CRY2	1,715 nondiabetic adults	Liver fat content Fasting glucose		Four <i>CRY2</i> SNPs were associated with fasting glucose — minor alleles were associated with elevated fasting glucose and reduced liver fat content <i>CRY2</i> mRNA expression from liver samples was associated with liver-fat content	(100)
CLOCK	897 adults aged 20–75 with coronary heart disease without clinical events in 6 months	Lipid metabolism HDL/ApoA1 ratio CRP	LFD, Mediterranean diet	<i>CLOCK</i> rs4580704 SNP C/C carriers had greater decrease in CRP and increase in HDL/ApoA1 following LFD compared with minor G allele carriers	(101)

CRP, C-reactive protein; LFD, low-fat diet.

across tissues. An important consequence of this definition is that many apparent risks can be misidentified or misdiagnosed. Take the “shift worker” designation. In one extreme, a subpopulation of shift workers may pay remarkable attention to circadian hygiene and may shift zeitgebers such as artificial light and mealtimes in accordance with their shifted schedule. In such a case, this subpopulation may not experience any circadian disruption despite being shift workers. The focus then becomes not just specifically modulation of sleep patterns or shift-working paradigms but, more importantly, those critical behavioral patterns, such as the timing of nutrient intake in alignment with the appropriate phases and amplitude of the internal clock (projected as the active phase).

Another critical consideration is the organ-specific function of the circadian clock and the distinct time to reentrainment of the clock behaviorally and in various organ systems following a circadian disruption. Internal organ reentrainment may not be mirrored by behavioral realignment, given the discordance identified between time to reentrainment in circadian behavioral patterns (activity/rest) and metabolic organ circadian rhythms (e.g., pancreas). This lends support to the value of cataloging predictably oscillating CCGs — or downstream clock-controlled genes — as useful biomarkers to determine an individual's internal circadian time. Such biomarkers would be invaluable in circadian-rhythm research, circadian-targeted drug dosing, clinical diagnostics, and epidemiology to identify at-risk populations. Various groups have undertaken this challenge, utilizing artificial intelligence and

machine learning applied to a range of biological sample types in both mice and humans, including whole blood (89–91), monocytes (92), hair-follicle cells (93), skin (94), breath (95), and multi-organ tissue sampling from autopsy (96).

In parallel, human studies offer mounting evidence linking circadian disruption to metabolic health, with a wide range of study types echoing and complementing each other's findings in analogous animal studies. However, existing studies are limited by the inherent nature of human subject research, lack of diversity in sample size, and lack of long-term studies in real-world settings. Nonetheless, phenotypic associations between numerous metabolic health outcomes and circadian disruptors, including circadian misalignment backed up by mechanistic animal studies, support the notion that CCGs are important systemic regulators in metabolic disorders. Future human research is needed to expand this evidence base and further illuminate the details of these relationships.

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1. Panda S, et al. Circadian rhythms from flies to human. *Nature*. 2002;417(6886):329–335.
2. Patke A, et al. Molecular mechanisms and physiological importance of circadian rhythms. *Nat Rev Mol Cell Biol*. 2020;21(2):67–84.
3. Chaix A, et al. The circadian coordination of cell biology. *J Cell Biol*. 2016;215(1):15–25.
4. Partch CL, et al. Molecular architecture of the mammalian circadian clock. *Trends Cell Biol*. 2014;24(2):90–99.
5. Yang X, et al. Nuclear receptor expression links the circadian clock to metabolism. *Cell*. 2006;126(4):801–810.
6. Hogenesch JB, et al. Characterization of a subset of the basic-helix-loop-helix-PAS superfamily that interacts with components of the dioxin signaling pathway. *J Biol Chem*. 1997;272(13):8581–8593.
7. Hogenesch JB, et al. The basic-helix-loop-helix-PAS orphan MOP3 forms transcriptionally active complexes with circadian and hypoxia factors. *Proc Natl Acad Sci U S A*. 1998;95(10):5474–5479.
8. Matsumura R, et al. The role of cell-autonomous circadian oscillation of Cry transcription in circadian rhythm generation. *Cell Reports*. 2022;39(3):110703.
9. Takahashi JS. Molecular components of the circadian clock in mammals. *Diabetes Obes Metab*. 2015;17 Suppl 1(0 1):6–11.
10. Acosta-Rodríguez VA, et al. Importance of circadian timing for aging and longevity. *Nat Commun*. 2021;12(1):2862.
11. Zhang R, et al. A circadian gene expression atlas in mammals: implications for biology and medicine. *Proc Natl Acad Sci U S A*. 2014;111(45):16219–16224.
12. Liu Y, et al. Prevalence of healthy sleep duration among adults—United States, 2014. *MMWR Morb Mortal Wkly Rep*. 2016;65(6):137–141.
13. Khubchandani J, Price JH. Short sleep duration in working American adults, 2010–2018. *J Community Health*. 2020;45(2):219–227.
14. Roenneberg T, et al. Social jetlag and obesity. *Curr Biol*. 2012;22(10):939–943.
15. Wittmann M, et al. Social jetlag: misalignment of biological and social time. *Chronobiol Int*. 2006;23(1-2):497–509.
16. Stokkan KA, et al. Entrainment of the circadian clock in the liver by feeding. *Science*. 2001;291(5503):490–493.
17. Chaix A, et al. Time-restricted feeding is a preventative and therapeutic intervention against diverse nutritional challenges. *Cell Metab*. 2014;20(6):991–1005.
18. Arble DM, et al. Circadian timing of food intake contributes to weight gain. *Obesity (Silver Spring)*. 2009;17(11):2100–2102.
19. Pickel L, Sung HK. Feeding rhythms and the circadian regulation of metabolism. *Front Nutr*. 2020;7:39.
20. Regmi P, Heilbronn LK. Time-restricted eating: benefits, mechanisms, and challenges in translation. *iScience*. 2020;23(6):101161.
21. Yoshida C, et al. Early nocturnal meal skipping alters the peripheral clock and increases lipogenesis in mice. *Nutr Metab (Lond)*. 2012;9(1):78.
22. Chaix A, et al. Time-restricted eating to prevent and manage chronic metabolic diseases. *Annu Rev Nutr*. 2019;39:291–315.
23. Salman EJ, Kabir R. Night eating syndrome. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing. 2024;NBK585047.
24. Daugaard S, et al. Light exposure during days with night, outdoor, and indoor work. *Ann Work Expo Health*. 2019;63(6):651–665.
25. Maury E. Off the clock: from circadian disruption to metabolic disease. *Int J Mol Sci*. 2019;20(7):1597.
26. Xie Y, et al. New insights into the circadian rhythm and its related diseases. *Front Physiol*. 2019;10:682.
27. Nakao A, et al. The circadian clock functions as a potent regulator of allergic reaction. *Allergy*. 2015;70(5):467–473.
28. Kiessling S, et al. Adrenal glucocorticoids have a key role in circadian resynchronization in a mouse model of jet lag. *J Clin Invest*. 2010;120(7):2600–2609.
29. Wever RA. Phase shifts of human circadian rhythms due to shifts of artificial Zeitgebers. *Chronobiologia*. 1980;7(3):303–327.
30. Schwartz PB, et al. Chronic jetlag alters the landscape of the pancreatic lipidome. *Pancreas*. 2022;51(1):80–89.
31. Nicholls SK, et al. Evidence for internal desynchrony caused by circadian clock resetting. *Yale J Biol Med*. 2019;92(2):259–270.
32. Alberti KG, et al. Harmonizing the metabolic syndrome: a joint interim statement of the International Diabetes Federation Task Force on Epidemiology and Prevention; National Heart, Lung, and Blood Institute; American Heart Association; World Heart Federation; International Atherosclerosis Society; and International Association for the Study of Obesity. *Circulation*. 2009;120(16):1640–1645.
33. Roenneberg T, Mellow M. The circadian clock and human health. *Curr Biol*. 2016;26(10):R432–R443.
34. Sun M, et al. Meta-analysis on shift work and risks of specific obesity types. *Obes Rev*. 2018;19(1):28–40.
35. Ward E. Carcinogenicity of night shift work. *Lancet Oncol*. 2019;20(8):1058–1059.
36. Rudic RD, et al. BMAL1 and CLOCK, two essential components of the circadian clock, are involved in glucose homeostasis. *PLoS Biol*. 2004;2(11):e377.
37. Turek FW, et al. Obesity and metabolic syndrome in circadian Clock mutant mice. *Science*. 2005;308(5724):1043–1045.
38. Sakhivel SJ, et al. A scoping review on the association between night eating syndrome and physical health, health-related quality of life, sleep and weight status in adults. *Nutrients*. 2023;15(12):2791.
39. Bass J, Takahashi JS. Circadian integration of metabolism and energetics. *Science*. 2010;330(6009):1349–1354.
40. Sehgal A. Physiology flies with time. *Cell*. 2017;171(6):1232–1235.
41. Hardin PE, et al. Feedback of the Drosophila period gene product on circadian cycling of its messenger RNA levels. *Nature*. 1990;343(6258):536–540.
42. Antoch MP, et al. Functional identification of the mouse circadian Clock gene by transgenic BAC rescue. *Cell*. 1997;89(4):655–667.
43. Bunker MK, et al. Mop3 is an essential component of the master circadian pacemaker in mammals. *Cell*. 2000;103(7):1009–1017.
44. Bur IM, et al. The circadian clock components CRY1 and CRY2 are necessary to sustain sex dimorphism in mouse liver metabolism. *J Biol Chem*. 2009;284(14):9066–9073.
45. Marcheva B, et al. Disruption of the clock components CLOCK and BMAL1 leads to hypoinsulinemia and diabetes. *Nature*. 2010;466(7306):627–631.
46. Cheon S, et al. Glucocorticoid-mediated Period2 induction delays the phase of circadian rhythm. *Nucleic Acids Res*. 2013;41(12):6161–6174.
47. Okano S, et al. Unusual circadian locomotor activity and pathophysiology in mutant CRY1 transgenic mice. *Neurosci Lett*. 2009;451(3):246–251.
48. Shan Z, et al. Rotating night shift work and adherence to unhealthy lifestyle in predicting risk of type 2 diabetes: results from two large US cohorts of female nurses. *BMJ*. 2018;363:k4641.
49. Hemmer A, et al. The effects of shift work on cardio-metabolic diseases and eating patterns. *Nutrients*. 2021;13(11):4178.
50. Antunes LC, et al. Obesity and shift work: chronobiological aspects. *Nutr Res Rev*. 2010;23(1):155–168.
51. van Amelsvoort LG, et al. Duration of shiftwork related to body mass index and waist to hip ratio. *Int J Obes Relat Metab Disord*. 1999;23(9):973–978.
52. Perelis M, et al. Pancreatic β cell enhancers regulate rhythmic transcription of genes controlling insulin secretion. *Science*. 2015;350(6261):aac4250.
53. Opperhuizen AL, et al. Rodent models to study the metabolic effects of shiftwork in humans. *Front Pharmacol*. 2015;6:50.
54. Yasumoto Y, et al. Short-term feeding at the wrong time is sufficient to desynchronize peripheral clocks and induce obesity with hyperphagia, physical inactivity and metabolic disorders in mice. *Metabolism*. 2016;65(5):714–727.
55. Izumo M, et al. Differential effects of light and feeding on circadian organization of peripheral clocks in a forebrain Bmal1 mutant. *Elife*. 2014;3:e04617.
56. Schwartz PB, et al. Chronic jetlag-induced alterations in pancreatic diurnal gene expression. *Physiol Genomics*. 2021;53(8):319–335.
57. Heyde I, Oster H. Differentiating external zeitgeber impact on peripheral circadian clock resetting. *Sci Rep*. 2019;9(1):20114.
58. Gale JE, et al. Disruption of circadian rhythms accelerates development of diabetes through pancreatic beta-cell loss and dysfunction. *J Biol Rhythms*. 2011;26(5):423–433.
59. Hepler C, et al. Time-restricted feeding mitigates obesity through adipocyte thermogenesis. *Science*. 2022;378(6617):276–284.
60. Asher G, Sassone-Corsi P. Time for food: the intimate interplay between nutrition, metabolism, and the circadian clock. *Cell*. 2015;161(1):84–92.
61. Hatori M, et al. Time-restricted feeding without reducing caloric intake prevents metabolic diseases in mice fed a high-fat diet. *Cell Metab*. 2012;15(6):848–860.
62. Ramin C, et al. Night shift work at specific age ranges and chronic disease risk factors. *Occup Environ Med*. 2015;72(2):100–107.
63. Givens ML, et al. Shiftwork, sleep habits, and metabolic disparities: results from the survey of the Health of Wisconsin. *Sleep Health*. 2015;1(2):115–120.
64. Schiavo-Cardozo D, et al. Appetite-regulating hormones from the upper gut: disrupted control of xenin and ghrelin in night workers. *Clin Endo-*

- crinol (Oxf)*. 2013;79(6):807–811.
65. Parsons MJ, et al. Social jetlag, obesity and metabolic disorder: investigation in a cohort study. *Int J Obes (Lond)*. 2015;39(5):842–848.
 66. Makarem N, et al. Evening chronotype is associated with poor cardiovascular health and adverse health behaviors in a diverse population of women. *Chronobiol Int*. 2020;37(5):673–685.
 67. Wong PM, et al. Social jetlag, chronotype, and cardiometabolic risk. *J Clin Endocrinol Metab*. 2015;100(12):4612–4620.
 68. Taylor BJ, et al. Bedtime variability and metabolic health in midlife women: The SWAN sleep study. *Sleep*. 2016;39(2):457–465.
 69. Kim M, et al. Objectively measured night-to-night sleep variations are associated with body composition in very elderly women. *J Sleep Res*. 2015;24(6):639–647.
 70. Buxton OM, et al. Adverse metabolic consequences in humans of prolonged sleep restriction combined with circadian disruption. *Sci Transl Med*. 2012;4(129):129ra43.
 71. Depner CM, et al. Ad libitum weekend recovery sleep fails to prevent metabolic dysregulation during a repeating pattern of insufficient sleep and weekend recovery sleep. *Curr Biol*. 2019;29(6):957–967.
 72. Moon S, et al. Beneficial effects of time-restricted eating on metabolic diseases: a systemic review and meta-analysis. *Nutrients*. 2020;12(5):1267.
 73. Chellappa SL, et al. Daytime eating prevents internal circadian misalignment and glucose intolerance in night work. *Sci Adv*. 2021;7(49):eabg9910.
 74. Resuehr D, et al. Shift work disrupts circadian regulation of the transcriptome in hospital nurses. *J Biol Rhythms*. 2019;34(2):167–177.
 75. Bracci M, et al. Rotating-shift nurses after a day off: peripheral clock gene expression, urinary melatonin, and serum 17- β -estradiol levels. *Scand J Work Environ Health*. 2014;40(3):295–304.
 76. Kervezee L, et al. Simulated night shift work induces circadian misalignment of the human peripheral blood mononuclear cell transcriptome. *Proc Natl Acad Sci U S A*. 2018;115(21):5540–5545.
 77. Archer SN, et al. Mistimed sleep disrupts circadian regulation of the human transcriptome. *Proc Natl Acad Sci U S A*. 2014;111(6):E682–E691.
 78. Möller-Levet CS, et al. Effects of insufficient sleep on circadian rhythmicity and expression amplitude of the human blood transcriptome. *Proc Natl Acad Sci U S A*. 2013;110(12):E1132–E1141.
 79. Vieira E G. Altered clock gene expression in obese visceral adipose tissue is associated with metabolic syndrome. *PLoS One*. 2014;9(11):e111678.
 80. Puig LS, et al. Influence of obesity, weight loss, and free fatty acids on skeletal muscle clock gene expression. *Am J Physiol Endocrinol Metab*. 2020;318(1):E1–E10.
 81. Pivovarov O, et al. Regulation of the clock gene expression in human adipose tissue by weight loss. *Int J Obes (Lond)*. 2016;40(6):899–906.
 82. Wefers J, et al. Circadian misalignment induces fatty acid metabolism gene profiles and compromises insulin sensitivity in human skeletal muscle. *Proc Natl Acad Sci U S A*. 2018;115(30):7789–7794.
 83. Cedernaes J, et al. Acute sleep loss induces tissue-specific epigenetic and transcriptional alterations to circadian clock genes in men. *J Clin Endocrinol Metab*. 2015;100(9):E1255–E1261.
 84. Jamshed H, et al. Early time-restricted feeding improves 24-hour glucose levels and affects markers of the circadian clock, aging, and autophagy in humans. *Nutrients*. 2019;11(6):1234.
 85. Lundell LS, et al. Time-restricted feeding alters lipid and amino acid metabolite rhythmicity without perturbing clock gene expression. *Nat Commun*. 2020;11(1):4643.
 86. Dashti HS, et al. Habitual sleep duration is associated with BMI and macronutrient intake and may be modified by CLOCK genetic variants. *Am J Clin Nutr*. 2015;101(1):135–143.
 87. Škrlec I, et al. The association between circadian clock gene polymorphisms and metabolic syndrome: a systematic review and meta-analysis. *Biology (Basel)*. 2021;11(1):20.
 88. Moreno CRC, et al. Are we ready to implement circadian hygiene interventions and programs? *Int J Environ Res Public Health*. 2022;19(24):16772.
 89. Braun R, et al. Universal method for robust detection of circadian state from gene expression. *Proc Natl Acad Sci U S A*. 2018;115(39):9247–9256.
 90. Hughey JJ. Machine learning identifies a compact gene set for monitoring the circadian clock in human blood. *Genome Med*. 2017;9(1):19.
 91. Laing EE, et al. Blood transcriptome based biomarkers for human circadian phase. *Elife*. 2017;6:e20214.
 92. Wittenbrink N, et al. High-accuracy determination of internal circadian time from a single blood sample. *J Clin Invest*. 2018;128(9):3826–3839.
 93. Akashi M, et al. Noninvasive method for assessing the human circadian clock using hair follicle cells. *Proc Natl Acad Sci U S A*. 2010;107(35):15643–15648.
 94. Wu G, et al. A single-sample circadian biomarker that performs across populations and platforms. *Genome Med*. 2020;12(73):820811.
 95. Martinez-Lozano Sinues P, et al. Circadian variation of the human metabolome captured by real-time breath analysis. *PLOS ONE*. 2015;9(12):e114422.
 96. Ruben MD, et al. A database of tissue-specific rhythmically expressed human genes has potential applications in circadian medicine. *Sci Transl Med*. 2018;10(458):eaat8806.
 97. Kumar V. *Biological Timekeeping: Clocks, Rhythms and Behaviour*. Springer; 2017.
 98. Lin E, et al. Effects of circadian clock genes and health-related behavior on metabolic syndrome in a Taiwanese population: Evidence from association and interaction analysis. *PLoS One*. 2017;12(3):e0173861.
 99. Corella D, et al. CLOCK gene variation is associated with incidence of type-2 diabetes and cardiovascular diseases in type-2 diabetic subjects: dietary modulation in the PREDIMED randomized trial. *Cardiovasc Diabetol*. 2016;15:4.
 100. Machicao F, et al. Glucose-raising polymorphisms in the human clock gene cryptochrome 2 (CRY2) affect hepatic lipid content. *PLoS One*. 2016;11(1):e0145563.
 101. Gomez-Delgado F, et al. Chronic consumption of a low-fat diet improves cardiometabolic risk factors according to the CLOCK gene in patients with coronary heart disease. *Mol Nutr Food Res*. 2015;59(12):2556–2564.