The 2017 Nobel Prize in Physiology and Medicine, awarded jointly to Jeffrey C. Hall, Michael Rosbash, and Michael W. Young for their discoveries of molecular mechanisms controlling the circadian rhythm, offered the opportunity to reflect on how clockwork mechanisms in the body affect the management of diabetes (1).

Circadian rhythmicity relates to all cells in the human body and to the microbiome that coexists within its confines (2,3). Clock genes control intracellular metabolic processes, whereas tissue activity in aggregate is responsive to central clock rhythms entrained by circadian light/dark cycles (Figure 1) (2,3). The circadian system regulates energy homeostasis for the body and is vulnerable to disruption through aberrant feeding/fasting cycles (2,4). Metabolic dysfunction induced by such chrono-disruptive feeding behavior may be corrected by time-restricted feeding in synchronicity with our master resident pacemaker in the suprachiasmatic nucleus of the hypothalamus responsive to immutable celestial cues (2,5–7).

The liver is the central organ involved in nutrient processing and energy metabolism (7,8). It receives amino acids, monosaccharides, and free fatty acids from the gut, along with other trace elements and vitamins, and proceeds to discharge its duties, fabricating tissue and generating energy (7). This process is regulated by hormonal and neurological cues from other organs, most importantly the pancreas, adrenal glands, and gut, and is very much dependent on appropriate timing (3,7,9). Feeding not entrained to innate circadian rhythmicity results in metabolic aberrations and gut dysfunction (3,10,11).
Metabolic aberrations are most frequently expressed as supraphysiologic glucose and triglyceride levels (3,7,10). These affect many organs and, importantly, contribute to anatomic changes in the liver, where inappropriate energy storage preponderates in the form of fatty infiltration (12).

Gut dysfunction relates to motility issues and microbiome disruption, resulting in maldigestion (13,14). The gut, with its trillions of resident microbes, is in constant communication regarding the size and composition of meals with organs throughout the body via hormonal and neuronal pathways (4). Digestive hyperactivity during rest alarms the organism at a time when restorative mechanisms are due to be operational as dictated by the sleep cycle.

The importance of time-restricted feeding for metabolic health was highlighted by early observations on the diurnal variation in oral glucose tolerance favoring the morning meal (15). Many more studies have since shown that circadian rhythmicity influences all aspects of digestion, including gastric pH, hepatic and renal blood flow, serum hormone levels (e.g., insulin and cortisol), and liver enzyme activity. These observations have served to establish the nascent field of chronopharmacology. More importantly, they have permitted us to appreciate the effects of nutrient input on the activity-rest cycle.

Nutrient sensing drives several peripheral clocks, including the liver, and may disrupt synchronization with the central pacemaker (16,17). Such circadian misalignment, driven by artificial light and nighttime work (16), has long-term implications expressed as impaired glucose tolerance, hyperlipidemia, weight gain, and hypertension, which we have come to collectively characterize as the metabolic syndrome. Re-entraining the peripheral clocks is now being recognized as the province of a new field in personalized medicine, namely that of “chrononutrition.” As such, the timing of meals is equally as important as their quality and quantity for health maintenance and longevity.

Our observations emphasize the importance of chrononutrition in people with diabetes because even insulin sensitivity has been shown to exhibit circadian rhythmicity (7). Glycemic responses to meals are exaggerated in these individuals, more so when the meal is timed to be consumed out of sync with the metabolic pacemaker (18). Accordingly, we now recognize optimal hormonal and hepatic function beginning in the early morning (4:00 a.m.) and fading in the early evening (4:00 p.m.) (Figures 2 and 3) (7). Review of glucose records—increasingly more abundant in the era of continuous glucose monitoring (CGM), which permits review of several months of detailed glucose profiling and follow-up care—confirms that evening lifestyle defines morning glycemia, thus setting the stage for the day’s profile. Late-night feeding and night-shift work introduce a “jet-lag” experience, which, when sustained, virtually guarantees metabolic dysregulation.

We have delivered this message to our patients consistently for many years, and more urgently now as CGM has served to underscore the level of control individuals have over their level of morning glycemia and, by extension, over the mean amplitude of their glycemic excursions (MAGE) throughout the day (19). This control is independent of any diabetes medication regimen.

Patient cases presented here in graphic form are representative of results obtained from our patients with diabetes ($n = 243$) using profiling of glucose meter results with Glooko (Glooko, Mountain View, Calif.) and/or CGM records using the Freestyle Libre Pro (Abbott, Alameda, Calif.) and Dexcom G4/G5 systems (Dexcom, San Diego, Calif.) in the past 2 years.

The patient whose glucose data are depicted in Figure 3 presented with type 1 diabetes. Panel A, a standard modal day CGM data report from midnight to midnight, does not highlight the relationship of dinner to the morning (fasting) blood glucose. Panel B showcases a modified CGM depiction conforming to the new 4:00 a.m.-to-4:00 p.m. time-restricted feeding paradigm and clearly demonstrates the effects of dinner on morning glycemia. Panel C illustrates the glycemic benefit of lifestyle modification when redis-

![FIGURE 2. Proposed paradigm shift in diabetes management redirecting focus to the rest period exemplified by the “stress-sleep” pillar. Patients and providers appreciate this concept by reviewing glycemic profiles as 4:00 p.m. to 4:00 p.m. depictions, as opposed to midnight-to-midnight. This graphic modification emphasizes the defining role of dinner in determining the next day’s fasting (morning) glucose level.](image-url)
morning euglycemia with increasing doses of insulin, frequently based on counting ill-timed carbohydrate deliveries, results in hypoglycemia. Patients appreciate that redistributing calories in this manner permits them to enjoy a wide variety of foods while maintaining weight control and improving diabetes control without increasing medication requirements. With simple adherence to the new paradigm, glycemic control without medication becomes feasible (Figures 4 and 5).

The patient whose data are depicted in Figure 4 has type 2 diabetes. In Panel A, the patient was on insulin with conventional meal plan emphasizing (late) dinner. Implementing the new 4:00 a.m. to 4:00 p.m. time-restricted feeding paradigm allowed the patient to maintain euglycemia off insulin, as shown in Panel B. This patient’s average fasting glucose before implementing the new guidelines was 103 mg/dL (25th–75th percentile 99–105 mg/dL) while on medication, including insulin. After implementing the new approach to meal timing, her average fasting glucose fasting was 112 mg/dL (25th–75th percentile 107–117 mg/dL) off of all medications.

Figure 5 provides similar depictions of glucose data from another patient with type 2 diabetes. This patient’s average fasting glucose before the meal timing restriction was 132 mg/dL (25th–75th percentile 119–143 mg/dL) while on medication. After implementing the new meal timing paradigm, the patient’s fasting glucose was 136 mg/dL (25th–75th percentile 123–144 mg/dL) without any diabetes medication.

Patients who modified their lifestyle to approximate the 4 a.m. to 4:00 p.m. feeding paradigm were the most successful in losing weight and improving their glycemic profile, with fewer medications. In all 243 patients providing records, the relationship of high fasting values with late meal times was evident.
Our intention in this article is to alert practitioners to the defining benefit of early meal times in achieving euglycemia. Insulin resistance can now be understood as a protective mechanism in an era of energy surfeit with inopportune deliveries. The liver, when challenged during its gut clock–dictated rest phase, stores and regurgitates rather than processes nutrients intended for energy needs and tissue growth and restoration (7,14). The benefits of fasting have been extolled, but the importance of timed fasting, according to gut clock dictates, remains to be reaffirmed and reintroduced to our lifestyle. The alternative, offered by polypharmacy, is both self-defeating and financially untenable for society (20).

**Duality of Interest**

No potential conflicts of interest relevant to this article were reported.

**Author Contributions**

N.H.E.M. wrote the manuscript, and V.B. researched the data and created the figures. Both authors reviewed and edited the manuscript. N.H.E.M. is the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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