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Human skeletal myotubes display a cell-autonomous circadian clock implicated in basal myokine secretion

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ABSTRACT

Objective: Circadian clocks are functional in all light-sensitive organisms, allowing an adaptation to the external world in anticipation of daily environmental changes. In view of the potential role of the skeletal muscle clock in the regulation of glucose metabolism, we aimed to characterize circadian rhythms in primary human skeletal myotubes and investigate their roles in myokine secretion.

Methods: We established a system for long-term bioluminescence recording in differentiated human myotubes, employing lentivector gene delivery of the *Bmal1-luciferase* and *Per2-luciferase* core clock reporters. Furthermore, we disrupted the circadian clock in skeletal muscle cells by transfecting siRNA targeting *CLOCK*. Next, we assessed the basal secretion of a large panel of myokines in a circadian manner in the presence or absence of a functional clock.

Results: Bioluminescence reporter assays revealed that human skeletal myotubes, synchronized *in vitro*, exhibit a self-sustained circadian rhythm, which was further confirmed by endogenous core clock transcript expression. Moreover, we demonstrate that the basal secretion of IL-6, IL-8 and MCP-1 by synchronized skeletal myotubes has a circadian profile. Importantly, the secretion of IL-6 and several additional myokines was strongly downregulated upon *siClock*-mediated clock disruption.

Conclusions: Our study provides for the first time evidence that primary human skeletal myotubes possess a high-amplitude cell-autonomous circadian clock, which could be attenuated. Furthermore, this oscillator plays an important role in the regulation of basal myokine secretion by skeletal myotubes.

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Keywords Circadian clock; Human skeletal myotube; Myokine; Interleukin-6; Circadian bioluminescence recording

1. INTRODUCTION

Circadian oscillations are daily cycles in behavior and physiology that have been described from photosynthetic bacteria to vertebrates. They are reflected by the existence of underlying intrinsic biological clocks with near 24 h periods that generate self-sustained rhythms, influenced by environmental stimuli, such as light and feeding [1]. Under homeostatic conditions, the clock acts as a driver of metabolic processes with remarkable tissue specificity that reflects the unique demand of each tissue [2]. In peripheral organs, a large number of key metabolic functions are subject to daily oscillations, such as carbohydrate and lipid metabolism by the liver, and xenobiotic detoxification by the liver, kidney and small intestine [2,3]. In rodents, the presence of peripheral circadian oscillators and their impact on gene expression

and organ function has been demonstrated in liver, whole pancreas, pancreatic islets, and in adipose tissue (reviewed in [4]). Of note, feeding rhythms represent a potent synchronizing cue for peripheral oscillators. Elegant studies involving inverted or restricted feeding schedules convincingly demonstrate that feeding rhythms are powerful enough to uncouple liver and other organ clocks from the SCN [5]. Moreover, rhythmicity in a number of clock-deficient mouse models could be restored by feeding rhythms [6–8].

There is growing evidence for a tight reciprocal link between a number of metabolic diseases, including obesity and type 2 diabetes mellitus (T2D), and the circadian clockwork [3,4]. Mice with circadian clock ablation develop hyperphagia, obesity, hyperglycemia and hypoinsulinemia [9]. The adipocyte-specific *Bmal1* knockout leads to obesity development [10], while the islet-specific *Bmal1* ablation

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Abbreviations

BMAL1	brain and muscle ARNT-like 1
CLOCK	circadian locomotor output cycles kaput
CRY	cryptochrome
DBP	D-albumin binding protein
GLP-1	glucagon-like peptide 1
HPRT	hypoxanthine-guanine phosphoribosyltransferase
IL-6	interleukin-6
IL-8	interleukin-8
Luc	luciferase
MOI	multiplicity of infection
MCP-1	monocyte chemotactic protein 1
M-CSF	macrophage colony-stimulating factor
PER	period
ROR α	retinoid-related orphan receptor alpha
REV-ERB α	reverse-erb alpha
SCN	suprachiasmatic nucleus
T2D	type 2 diabetes mellitus
VEGF	vascular endothelial growth factor
ZT	zeitgeber time

directly triggers the onset of T2D in mice [11]. In humans, genetic analyses have shown a close association between glucose levels and variants of the *CRY2* and melatonin receptor 1B (*MTNR1B*) genes [2]. Skeletal muscle is the largest insulin sensitive organ in the body, playing an essential role in whole body glucose homeostasis. It is responsible for 70–80% of insulin-stimulated glucose uptake and therefore representing the most important site of insulin resistance in T2D patients [12]. The primary and best-described function of skeletal muscles is their mechanical activity. During the last decade, however, skeletal muscle has been characterized also as a secretory organ, producing and releasing myokines that act on the muscle itself and display endocrine effects on distant organs [13,14]. Recently, it has been proposed that fully differentiated primary human skeletal muscle cells secrete over 300 potential myokines [15]. Interleukin-6 (IL-6) is one of the first identified myokines, which is produced by the contracting skeletal muscle [16]. Acute high plasma levels of IL-6 are associated with exercise, while chronically elevated IL-6 is observed upon obesity and T2D. Moreover, IL-6 promotes pancreatic alpha cell mass expansion [17] and stimulates GLP-1 production and secretion by alpha and L-cells upon metabolic syndrome, thus exerting beneficial effects in T2D mouse models [18]. Besides IL-6, skeletal muscle produces a number of additional myokines, such as MCP-1 and IL-8, which play a role in skeletal muscle inflammation, recruitment of macrophages and insulin sensitivity [19,20].

In rodents, about 7% of the skeletal muscle transcriptome is expressed in a circadian manner [21]. Moreover, clock ablation (*Bmal1*^{-/-}) leads to skeletal muscle pathologies [22]. *MyoD*, a master regulator of myogenesis, exhibits a robust circadian rhythm and was identified as a direct target of CLOCK and BMAL1. In addition, disruption of myofilament organization was detected in *Bmal1*^{-/-}, *Clock* ^{Δ 19}, and in *MyoD*^{-/-} mice, suggesting a direct link between the circadian clock and skeletal muscle function in rodents [22]. Furthermore, a recent study suggests that in mice muscle insulin sensitivity might be clock-dependent [23].

In view of the accumulating evidence in rodent models, it has been accepted that the skeletal muscle clock plays an essential role in maintaining proper metabolic functioning (reviewed in [24]), although the mechanism of this important regulation is not entirely clear,

particularly in human subjects. Given that, the circadian clock in human skeletal muscle has remained largely unexplored, we aimed to characterize the circadian oscillator in primary human myotubes and explore its impact on the regulation of human skeletal muscle myokine secretion.

2. MATERIAL AND METHODS

2.1. Study participants, skeletal muscle tissue sampling and primary cell culture

Muscle biopsies were derived from non-obese or obese donors with the informed consent obtained from all participants (see Table 1 for the

Table 1 — Characteristics of donors for skeletal muscle biopsies.

Donor	Sex	Age (years)	BMI (kg/m ²)	Biopsy source
Non-obese	M = 14, F = 5	58 ± 18	24.88 ± 3.20	
1 ^a	M	48	21.7	Rectus abdominus
2 ^{a,f}	M	45	21	Rectus abdominus
3 ^{a,f}	F	58	20.5	Rectus abdominus
4 ^{a,f}	F	42	19.5	Rectus abdominus
5 ^{b,d,k}	M	23	29.34	Gluteus maximus
6 ^{a,b,d,e,g,h,i,j}	M	62	24.3	Gluteus maximus
7 ^{a,b,f}	F	77	25.6	Gluteus maximus
8 ^{b,d,g,i,j}	M	57	26	Gluteus maximus
9 ^{b,d,g,h}	M	60	24	Gluteus maximus
10 ^b	F	88	29.64	Gluteus maximus
11 ^{a,b,c,d}	F	65	25.8	Gluteus maximus
12 ^{b,d,k}	M	25	19.27	Gluteus maximus
13 ^{b,d,g,h}	M	64	28	Gluteus maximus
14 ^{b,d}	M	87	25.51	Gluteus maximus
15 ^{b,d}	M	85	25.54	Gluteus maximus
16 ^{b,d}	M	72	26.5	Gluteus maximus
17 ^{b,e,f,h}	M	48	24.3	Gluteus maximus
18 ^{b,d,h}	M	45	28.1	Gluteus maximus
19 ^{a,b,c,d,g,i,j}	M	57	28.1	Gluteus maximus
Obese	M = 1, F = 4	53 ± 15	39.21 ± 7.16	
20 ^b	M	70	30.1	Gluteus maximus
21 ^e	F	70	32.9	Gluteus maximus
22 ^{b,e,g,h,i}	F	43	43	Gluteus maximus
23 ^{b,d,g,j}	F	39	45.48	Rectus abdominus
24 ^{b,d,g,i}	F	45	44.58	Rectus abdominus
All donors	M = 15, F = 9	57 ± 17	27.87 ± 7.23	

M, male; F, female.

Non-obese, data are mean ± SD, *n* = 19.

Obese, data are mean ± SD, *n* = 5.

All donors, data are mean ± SD, *n* = 24.

^a donor cells used for the recording of *Bmal1-luc* bioluminescence of dexamethasone-synchronized samples.

^b donor cells used for the recording of *Bmal1-luc* bioluminescence of forskolin-synchronized samples.

^c donor cells used for the recording of *Bmal1-luc* bioluminescence of dexamethasone vs. forskolin-synchronized samples.

^d donor cells used for the recording of *Per2-luc* bioluminescence of forskolin-synchronized samples.

^e donor cells used for the around-the-clock experiment with dexamethasone synchronization.

^f donor cells used to quantify the silencing of *CLOCK* in *siControl* / *siClock* samples synchronized with dexamethasone.

^g donor cells used to quantify the silencing of *CLOCK* in *siControl* / *siClock* samples synchronized with forskolin.

^h donor cells used for the around-the-clock experiment in forskolin-synchronized *siControl* / *siClock* samples.

ⁱ donor cells used for the IL-6 perfusion experiments with forskolin synchronization.

^j donor cells used for the multiplex assay analysis of perfusion samples synchronized with forskolin.

^k donor cells used for the IL-6 perfusion experiments with forskolin vs. dexamethasone synchronization.

donor characteristics). The experimental protocol ('DIOMEDE') was approved by the Ethical Committee SUD EST IV (Agreement 12/111) and performed according to the French legislation (Huriet's law). All donors had HbA_{1c} levels inferior to 6.0% and fasting glycemia inferior to 7 mmol/L, were not diagnosed for T2D, neoplasia or chronic inflammatory diseases, and not doing shift work. Biopsies were taken from the *Gluteus maximus* ($n = 18$) or the *Rectus abdominus* ($n = 6$) muscles during the planned surgeries. Primary skeletal myoblasts were purified and differentiated into myotubes according to the previously described procedure [25]. Briefly, after removal of apparent connective and fat tissue contaminants, the muscle biopsy was minced with scissors and incubated successively at least 4 times for 20 min in Trypsin-EDTA (Invitrogen, Thermo Fisher Scientific, Waltham, MA, USA) at 37 °C under agitation. Trypsin-EDTA digested extracts were pooled, and centrifuged (150 g). The pellet was rinsed several times with PBS and cells were filtered on a 70- μ m filter before being plated in a Primaria flask (Falcon; Becton Dickinson, Bedford, MA) containing growth medium composed of HAM F-10 supplemented with 2% Ultrosor G (BioSeptra SA, Cergy-Saint-Christophe, France), 2% fetal bovine serum (FBS, Invitrogen), and 1% antibiotics (Invitrogen). After 4 days, the myoblasts were immuno-selected using a monoclonal human CD56 antibody combined with paramagnetic beads (CD56 MicroBeads, Miltenyi Biotec, Germany), according to the manufacturer's instructions. The selected cells were plated on Primaria plastic ware at 4500 cells/cm² and cultured in growth medium at 37 °C. After reaching confluence, myoblasts were differentiated into myotubes during 7–10 days in DMEM supplemented with 2% FBS. Muscle differentiation was characterized by the fusion of myoblasts into polynucleated myotubes (Supplementary Figure 1).

2.2. siRNA transfection and lentiviral transduction

Human primary myoblasts were differentiated into myotubes as described above. Cells were transfected with 20 nM siRNA targeting *CLOCK* (*siClock*), or with non-targeting *siControl* (Dharmacon, GE Healthcare, Little Chalfont, UK), using HiPerFect transfection reagent (Qiagen, Hombrechtikon, Switzerland) following the manufacturer's protocol, 24 h prior to synchronization. To produce lentiviral particles, *Bmal1-luc* [26] or *Per2-luc* [27] lentivectors were transfected into 293T cells using the polyethylenimine method (for detailed procedure see [28]). Myoblasts were transduced with the indicated lentiviral particles with a multiplicity of infection (MOI) = 3 for each, grown to confluence, and subsequently differentiated into myotubes.

2.3. In vitro skeletal myotube synchronization and real-time bioluminescence recording

To synchronize primary myotubes, 10 μ M forskolin (Sigma, Saint-Louis, MO, USA) or 100 nM dexamethasone (Alfa Aesar, Johnson Matthey, London, UK) were added to the culture medium, respectively. Following 60 min (forskolin) or 30 min (dexamethasone) incubation at 37 °C in a cell culture incubator, the medium was changed to a phenol red-free recording medium containing 100 μ M luciferin and cells were transferred to a 37 °C light-tight incubator (Prolume LTD, Pinetop, AZ, USA), as previously described by us [28]. Bioluminescence from each dish was continuously monitored using a Hamamatsu photomultiplier tube (PMT) detector assembly. Photon counts were integrated over 1 min intervals. Luminescence traces are either shown as raw or detrended data. For detrended time series, luminescence signals were smoothed by a moving average with a window of 144 data points and detrended by an additional moving average with a window of 24 h [29]. For quantification of the circadian amplitude and period the first cycle was not taken into consideration.

2.4. mRNA extraction and quantitative PCR analysis

Differentiated myotubes were synchronized by forskolin or dexamethasone, collected every 4 h during 48 h (0 h–48 h), or during 24 h (12 h–36 h), deep-frozen in liquid nitrogen and kept at –80 °C. Total RNA was prepared using RNeasy Plus Micro kit (Qiagen). 0.5 μ g of total RNA was reverse-transcribed using Superscript III reverse transcriptase (Invitrogen) and random hexamers and PCR-amplified on a LightCycler 480 (Roche Diagnostics AG, Rotkreuz, Switzerland). Mean values for each sample were calculated from the technical duplicates of each qRT-PCR analysis, and normalized to the mean of two housekeeping genes (*HPRT* and *9S* or *GAPDH* and *9S*), which served as internal controls. Primers used for this study are listed in Supplementary Table 1.

2.5. Circadian analysis of basal myokine secretion by ELISA and multiplex assay

In vitro synchronized differentiated myotubes, transduced with *Bmal1-luc* reporter, were placed into an in-house developed two-well horizontal perfusion chamber, connected to the LumiCycle. Cells were continuously perfused for 48 h with culture medium containing 100 μ M luciferin. Bioluminescence recordings were performed in parallel to the automated collection of outflow medium in 4 h intervals. Basal IL-6 levels were quantified in the outflow medium using the Human IL-6 Instant ELISA kit (eBioscience, Affymetrix, Santa Clara, CA, USA) following the manufacturer's instructions. Data were normalized to the genomic DNA content, extracted using the QIAamp DNA Blood Mini Kit (Qiagen). Perfusion medium samples were further concentrated using Amicon Ultra 2 ml centrifugal filters (Ultracel-3K, Merck Millipore, Darmstadt, Germany). The evaluation of myokine release from human primary skeletal muscle cells was carried out using a multiplex bead array assay system (R&D Systems, Minneapolis, MN, USA). Custom-made luminex screening plates (CD44, CHI3L1, FABP3, galectin-3, GRO- α , IGFBP-3, IL-7, IL-13, IL-17A, M-CSF, MCP-1, MMP-2, Serpin C1, Serpin E1, TIMP-1) and high sensitivity performance plates (IL-1 beta, IL-2, IL-4, IL-6, IL-8, IL-10, IL-12 p70, IFN- γ , TNF- α and VEGF) were analyzed according to the manufacturer's instructions. Plate analysis was performed on a Bio-Plex 200 array reader (Bio-Rad Laboratories, Hercules, CA, USA), with the Bio-Plex software (Bio-Rad) used for data analysis.

2.6. Data analysis

Actimetrics LumiCycle analysis software (Actimetrics LTD) and the JTK_CYCLE algorithm [30] were used for bioluminescence and myokine secretion profile data analyses, respectively. For the ELISA and multiplex data analysis, 2 technical duplicates from 3 biological samples were analyzed for each myokine. From these values the average \pm the SEM was calculated for each time point. For JTK_CYCLE analysis the period width was set at 20–24 h. Statistical analyses were performed using a paired Student's *t*-test. Differences were considered significant for $p < 0.05$ (*), $p < 0.01$ (**), and $p < 0.001$ (***)

3. RESULTS

3.1. High-amplitude self-sustained clocks are functional in primary human skeletal myotubes

Circadian bioluminescence recordings in living cells allow for the study of molecular clocks in human peripheral tissues, as previously demonstrated by us [28,31] and others [32]. We applied this powerful methodology to assess clock properties in human primary skeletal myotubes established from human donor biopsies and differentiated *in vitro* (see Supplementary Figure 1 and Table 1 for donor

characteristics). Multiple *in vitro* stimuli have previously been demonstrated to efficiently synchronize cultured cells, among them forskolin and dexamethasone [28,33]. As shown in Supplementary Figure 2, short pulses of dexamethasone or forskolin were able to strongly synchronize human myotubes bearing the *Bmal1-luc* lentivector. Both forskolin and dexamethasone induced oscillations with comparable period length: 25.29 ± 0.13 h, $n=19$ for forskolin (Table 2), and 24.48 ± 0.24 h, $n=8$ for dexamethasone (data not shown). However, the forskolin pulse induced more sustained cycles with higher circadian amplitude compared to dexamethasone-synchronized cells (Supplementary Figure 2) and was therefore mainly used in this study as the *in vitro* synchronization stimulus. The here established experimental settings allowed for continuous recording of oscillation profiles in human primary myotubes for several days with high resolution (Figure 1A, B). As expected, the profiles of the *Bmal1-luc* and *Per2-luc* reporters were antiphasic (Figure 1A, B; [3]). High-amplitude self-sustained oscillations were reproducibly recorded from human primary myotubes for *Bmal1-luc* and *Per2-luc* reporters with an average period length of 25.29 ± 0.13 h and 25.20 ± 0.19 h, respectively (Figure 1C). Of note, no significant difference in period length of the *Bmal1-luc* or *Per2-luc* reporter was observed between myotubes established from non-obese and obese donors (Table 2), which might reflect the resistance of the core clock to metabolic changes.

To further validate the results obtained by circadian bioluminescence reporter studies, we examined endogenous core clock gene expression profiles in forskolin-synchronized myotubes (Figure 2C, closed circles). mRNA accumulation patterns from synchronized skeletal myotubes were monitored every 4 h during 48 h by quantitative RT-PCR, using amplicons for *BMAL1*, *REV-ERB α* , *PER3* and *DBP*. The values obtained were normalized to the mean of the housekeeping genes *HPRT* and *9S*. In agreement with our *Bmal1-luc* reporter experiments, endogenous *BMAL1* transcript levels exhibited circadian oscillations over 48 h in synchronized myotubes (compare Figure 2C *BMAL1* panel to Figure 1A), clearly antiphasic to the profiles of the endogenous *REV-ERB α* , *PER3*, and *DBP* transcripts (Figure 2C). Similar experiments were conducted in dexamethasone-synchronized myotubes (Supplementary Figure 3). *BMAL1* and *CRY1* mRNAs exhibited oscillatory profiles antiphasic to those of *REV-ERB α* and *PER2-3*, consistent with the dexamethasone-induced oscillations of the *Bmal1-luc* reporter (Supplementary Figure 2).

3.2. Human myotube clock disruption by siRNA-mediated *CLOCK* knockdown

In order to disrupt the circadian clock in cultured human myotubes, we set up an efficient *siClock* transfection protocol, resulting in more than 80% knockdown of *CLOCK* transcript levels (Figure 2A, Supplementary Figure 4A). Circadian expression of the *Bmal1-luc* reporter was blunted in *siClock*-transfected myotubes upon forskolin or dexamethasone synchronization, if compared to cells transfected with non-targeting

sequences (*siControl*) or to non-transfected counterparts (Figure 2B, Supplementary Figure 4B), thus validating circadian clock disruption. Moreover, the amplitudes of endogenous *REV-ERB α* , *PER3* and *DBP* transcript profiles were strongly blunted in *siClock*-transfected cells, in comparison to *siControl* cells (Figure 2C, Supplementary Table 2). By contrast, *BMAL1* transcript levels were slightly upregulated (Figure 2C, Supplementary Table 2).

3.3. Regulation of basal IL-6 secretion by the circadian clock in human primary myotubes

Given the accumulating evidence on the essential role of IL-6 secretion by skeletal muscle under physiological conditions and upon metabolic diseases [14,17,18], we next monitored basal circadian IL-6 secretion by human primary myotubes established from non-obese and obese donors. To this end, we developed an in-house perfusion system connected to the LumiCycle chamber that allows for parallel cell perfusion and bioluminescence profile recordings. Basal IL-6 secretion from *in vitro* synchronized skeletal myotubes was monitored in “around-the-clock” experimental settings, with a continuous flow of culture medium (see Material and methods for details). The perfusion experiments suggested that forskolin-synchronized myotubes exhibited a circadian profile of basal IL-6 secretion over 48 h, with a *Zenith* around 8 h–12 h and 32 h–36 h and a *Nadir* of 20 h–24 h (Figure 3A). JTK_CYCLE analysis [30] revealed that average profile of basal IL-6 secretion did not reach significance to be qualified as circadian over the entire time span of 48 h. However it was significantly circadian over the first 36 h of perfusion following *in vitro* synchronization (** $p < 0.01$, $n = 6$). Similar basal IL-6 secretion profiles were observed from dexamethasone-synchronized myotubes (Supplementary Figure 5).

We next tested the effect of *CLOCK* depletion on IL-6 secretion. Similar to previous experiments (Figure 2), *CLOCK* transcript expression was at least 80% downregulated in *siClock*-transfected myotubes (** $p < 0.001$, paired *t*-test, Table 3) compared to *siControl*-transfected cells. The achieved clock disruption was also confirmed by parallel *Bmal1-luc* bioluminescence recording in perfused cells (Figure 3B). Importantly, IL-6 secretion decreased on average 64% in *siClock*-transfected myotubes compared to *siControl*-transfected counterparts ($p < 0.05$, Table 3). Furthermore, the profile of basal IL-6 secretion became flat upon clock disruption if compared to the *siControl*-transfected profile (see red line vs. black line, Figure 3A). Of note, overall basal IL-6 secretion was on average higher in obese *siControl*-transfected subjects ($n = 3$) compared to non-obese *siControl*-expressing counterparts ($n = 3$), although values did not reach statistical significance (Table 3).

To get an insight into the regulation of basal IL-6 secretion by the circadian clock, we assessed *IL6* transcript levels following *in vitro* synchronization with forskolin. No clear circadian pattern was observed (Figure 3C). The strong immediate early peak of *IL6* transcription induced by the forskolin pulse might be attributed to the presence of a cAMP response element previously identified in IL-6 promoter region [34]. Moreover, *CLOCK* depletion by *siClock* had no effect on basal *IL6* transcription (Figure 3C).

3.4. Multiplex screen identifies additional clock-regulated myokines

Since we obtained convincing evidence on the requirement of a functional circadian clock for basal IL-6 secretion by skeletal myotubes (Figure 3A,B), we selected an additional panel of myokines for analysis by multiplex assay. The around-the-clock secretory profiles of IL-6 and 24 other myokines were assessed in perfusion samples obtained from

Table 2 — Circadian period length of forskolin-synchronized human myotubes assessed by circadian bioluminescent reporters.

	<i>Bmal1-luc</i> period (h)		<i>Per2-luc</i> period (h)		Mean period (h)	
	<i>n</i>	Mean \pm SEM	<i>n</i>	Mean \pm SEM	<i>n</i> *	Mean \pm SEM
Non-obese	15	25.35 \pm 0.14	12	25.25 \pm 0.22	27	25.31 \pm 0.12
Obese	4	25.04 \pm 0.35	2	24.85 \pm 0.30	6	24.98 \pm 0.24
All donors	19	25.29 \pm 0.13	14	25.20 \pm 0.19	33	25.25 \pm 0.11

**n* represents the number of experimental repetitions.

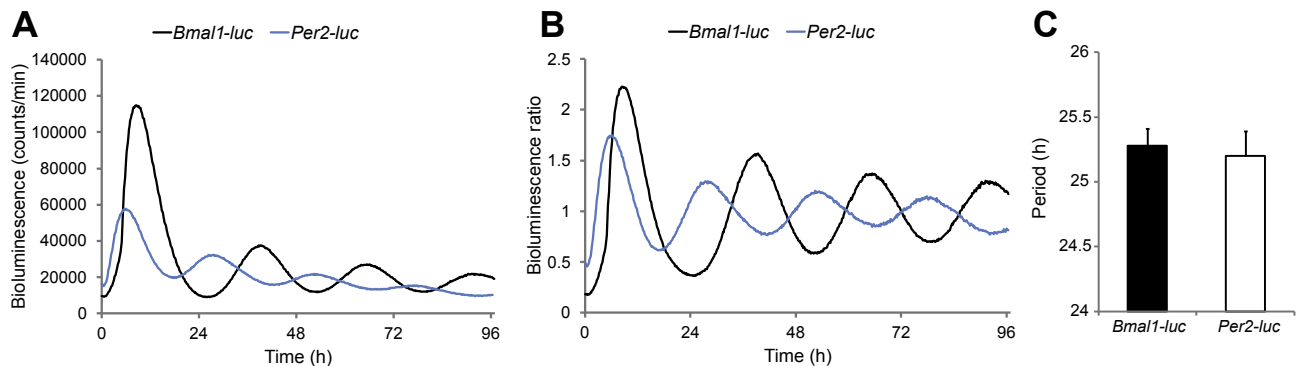


Figure 1: High-amplitude cell autonomous oscillators are functional in differentiated human primary myotubes. Human primary myoblasts were transduced with lentiviral particles expressing *Bmal1-luc* (black line) or *Per2-luc* (blue line). Cells were differentiated into myotubes, synchronized with forskolin, and transferred to the Actimetrics LumiCycle for bioluminescence recording. Raw (A) and detrended (B) oscillation profiles are representative of 19 and 14 independent experiments, respectively (one donor per experiment). (C) The period length of *Bmal1-luc* or *Per2-luc* was on average 25.29 ± 0.13 h, ($n = 19$) or 25.20 ± 0.19 h, ($n = 14$), respectively. Data represent the mean \pm SEM.

primary human myotubes as described in Figure 3, and further concentrated to allow for the detection of basal myokine secretion in *siControl* and *siClock*-transfected myotubes. Selected myokines were chosen based on their presence in the secretome of differentiated primary skeletal muscle cells [15], their function and implication in metabolic diseases, and their availability and compatibility for multiplex array analysis. Out of the selected panel of 25 myokines, 15 myokines were detected in the concentrated perfusion samples, while 10 myokines remained undetectable, due to their low levels of basal secretion (Table 4).

Importantly, the profile and concentration levels of around-the-clock IL-6 secretion measured by multiplex analysis were very similar to those obtained by ELISA (compare Figure 3D–A). JTK_CYCLE analysis [30] confirmed that the average profile of secreted IL-6 measured by multiplex analysis was significantly circadian within 48 h (Figure 3D, Table 4). In addition to IL-6, MCP-1 and IL-8 were secreted in a circadian manner according to JTK_CYCLE analysis (Figure 4A, Table 4). M-CSF and GRO- α exhibited a secretion profile that might be clock-controlled; however, the adjusted minimal p -value did not reach significance according to JTK_CYCLE analysis (Figure 4B, Table 4). Furthermore, although the temporal secretion patterns of VEGF, CD44, FABP3, Galectin-3 and TIMP-1 were not identified as circadian, the overall secretion levels of these myokines were significantly reduced upon *CLOCK* depletion (Figure 4C, Table 4).

4. DISCUSSION

4.1. Molecular makeup of the circadian oscillator operative in human skeletal muscle

Our study provides for the first time evidence for cell-autonomous self-sustained circadian oscillators, operative in human primary skeletal myotubes. Molecular characteristics of the human myotube clock were assessed by two complementary approaches. Pronounced circadian oscillations were recorded with high temporal resolution for at least 4–5 consecutive days from *in vitro* synchronized primary skeletal myotubes, transduced with *Bmal1-luciferase* or *Per2-luciferase* lentivectors (Figure 1A,B). The circadian characteristics of human skeletal myotubes were in accordance with those reported for human primary thyrocytes, pancreatic islets, skin fibroblasts [28,31,35], and for mouse skeletal muscle assessed *in vivo* [21,22]. Sustained circadian oscillations were efficiently induced in our system by both forskolin (Figure 1) and dexamethasone

(Supplementary Figure 2) pulses, suggesting that these oscillators are functional irrespective of the synchronization stimulus or entrainment pathway. It might be of interest to further explore whether other physiologically relevant stimuli like glucose, insulin, or myokine-induced signaling pathways play a role in human myotube synchronization.

In line with the outcome of our reporter experiments, endogenous around-the-clock gene expression analyses suggested that the core clock genes *BMAL1*, *REV-ERB α* , and *PER3*, as well as the clock output gene *DBP*, exhibit circadian oscillatory patterns in forskolin and dexamethasone-synchronized myotubes (Figure 2C closed circles, Supplementary Figure 3). Of note, the circadian amplitude of oscillations induced *in vitro* is typically lower if compared to *in vivo* oscillations from the same tissue, as demonstrated, for instance, for mouse islets synchronized *in vitro* or collected *in vivo* [11]. Thus, identifying clock-controlled genes *in vitro* by RT-qPCR represents a significant challenge due to rather low amplitudes [21,28]. More accurate methods like RNA sequencing of samples collected with higher temporal resolution might represent a solution to this problem.

4.2. Experimental model for circadian clock disruption in human primary myotubes

Here, we have established experimental settings for a highly reproducible siRNA-mediated *CLOCK* transcript knockdown of more than 80% in human primary muscle cells (Figure 2A, Supplementary Figure 4A). Upon such *CLOCK* silencing, significant flattening of the circadian amplitude was observed for the *Bmal1-luc* reporter (Figure 2B, Figure 3B, Supplementary Figure 4B) and for endogenous *REV-ERB α* , *PER3* and *DBP* transcripts (Figure 2C), confirming circadian core clock disruption. The discrepancy between *Bmal1-luc* reporter data and endogenous *BMAL1* expression upon *siClock* might be related to the fact that the knockdown effect of *CLOCK* on the *Bmal1-luc* reporter is evident primarily after 48 h, and that the promoter length of the *Bmal1* reporter is different from the endogenous gene. While *REV-ERB α* , *PER3* and *DBP* transcript profiles only exhibited residual circadian oscillations, which might be explained by incomplete clock ablation, the amplitude of the endogenous *BMAL1* oscillatory profile was not reduced and even slightly increased (Figure 2C). Similarly, siRNA-mediated depletion of *CLOCK* in U2OS cells was reported to increase *BMAL1* expression levels [36]. Moreover, *Clock*-deficient mice continue to exhibit circadian patterns of behavioral and molecular

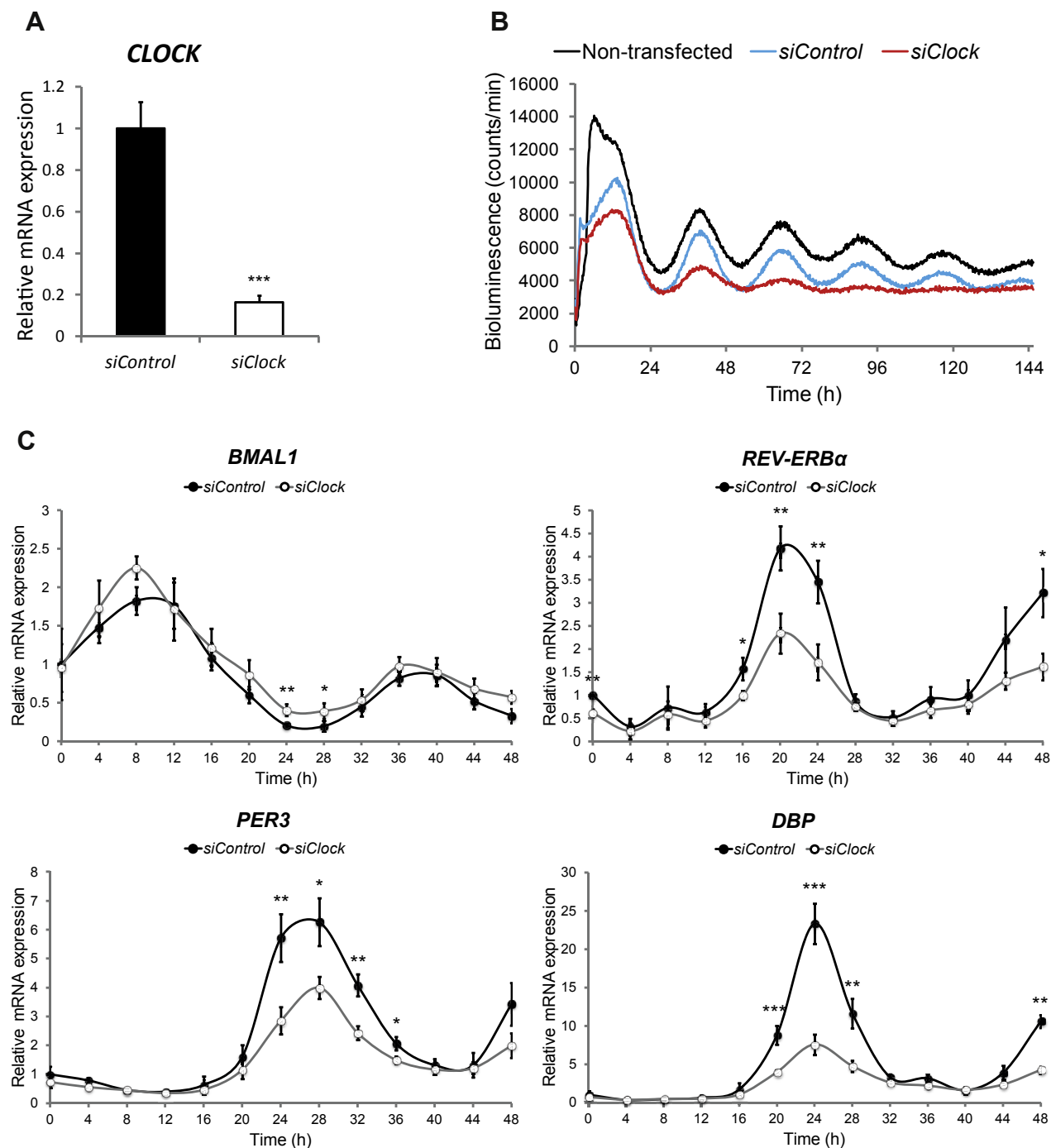


Figure 2: Silencing of *CLOCK* expression attenuates circadian oscillations in human skeletal myotubes. (A) *CLOCK* mRNA was measured in human myotubes transfected with *siControl* or *siClock* by RT-qPCR and normalized to the mean of *9S* and *HPRT*. *CLOCK* expression was reduced by $83.5 \pm 3.4\%$ (mean \pm SEM, $n = 8$; $***p < 0.001$) in *siClock*-transfected cells. (B) Amplitude of the *Bmal1-luc* reporter is strongly reduced in *siClock*-transfected myotubes. Representative *Bmal1-luc* oscillation profiles are shown for non-transfected (black line), *siControl* (blue line), and *siClock* (red line) transfected myotubes. *Bmal1-luc* oscillation profiles were recorded in duplicates (3 experiments, one donor per experiment). (C) RT-qPCR was performed for *BMAL1*, *REV-ERB α* , *PER3* and *DBP* on RNA samples extracted from forskolin-synchronized human myotubes, transfected with *siClock* (open circles) or *siControl* (closed circles). Samples were collected every 4 h and normalized to the mean of *9S/HPRT*. Profiles (mean \pm SEM) are representative of 3 experiments (2 donors for time points 0 h–48 h and 3 donors for time points 12 h–36 h) with duplicates per time point.

rhythms. In the SCN and liver of *Clock*-deficient mice, *Bmal1* mRNA was elevated, as well as in pancreatic islets of *Clock*-mutant mice, which was attributed to reduced REV-ERB α expression and a compensatory effect of NPAS2 [11,37].

Clockwork perturbations may develop in humans with ageing and upon a number of disorders [3,4]. In this respect, our experimental model, which allows for reproducible clock disruption mediated by *CLOCK* depletion in differentiated skeletal myotubes, represents a valuable

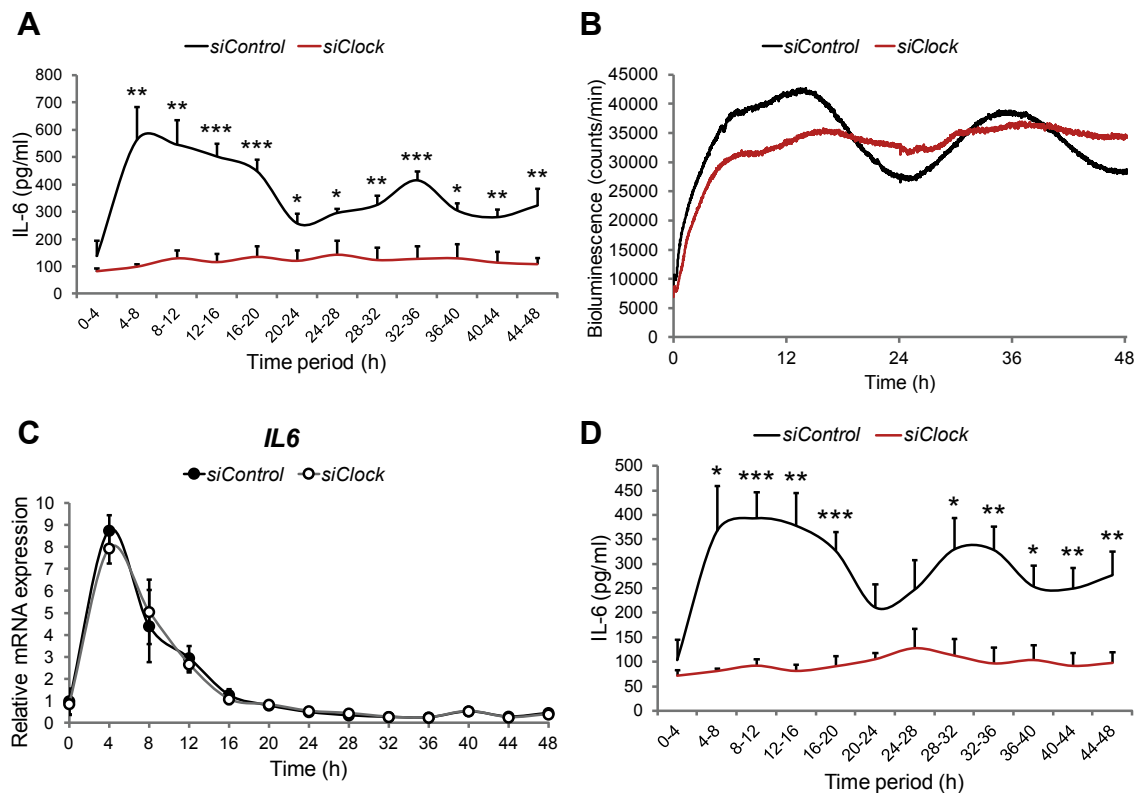


Figure 3: Basal IL-6 secretion by human skeletal myotubes is strongly inhibited in the absence of a functional circadian clock. Myoblasts were transduced with the *Bmal1-luc* lentivector, differentiated into myotubes and transfected with either *siControl* or *siClock* siRNA. 24 h following transfection, myotubes were synchronized with forskolin and subjected to continuous perfusion with parallel bioluminescence recording. (A) Basal IL-6 secretion profile (mean + SEM) in the presence or absence of a functional clock. The perfusion outflow medium was collected continuously in an automated manner in 4 h intervals until 48 h (0–4 corresponds to the accumulation of IL-6 between 0 h and 4 h). IL-6 levels in the perfusion outflow medium were assessed by ELISA. The results represent basal IL-6 levels normalized to the total DNA content. 2 technical duplicates from 3 independent experiments (3 non-obese donors, see Table 3) were analyzed for each time point. (B) *Bmal1-luc* bioluminescence profiles of *siControl*-transfected myotubes (black line) and *siClock*-transfected myotubes (red line), representative of 3 experiments, with one donor cell line used per experiment. (C) RT-qPCR was performed for *IL6* on RNA samples extracted from forskolin-synchronized human myotubes, transfected with *siClock* (open circles) or *siControl* (closed circles). Samples were collected every 4 h and normalized to the mean of *9S/HPRT*. Profiles (mean ± SEM) are representative of 3 experiments (2 donors for time points 0 h–48 h and 3 donors for time points 12 h–36 h) with duplicates per time point. (D) Basal IL-6 secretion profiles in the presence or absence of a functional clock obtained from concentrated perfusion samples, assessed by multiplex analysis. Data shown as mean + SEM of 2 technical duplicates from 3 biological samples for each time point, normalized to the total DNA content.

tool for getting significant insights into the roles of clock perturbations in different aspects of human skeletal muscle function. In order to detect oscillatory alterations caused by metabolic changes it might be necessary to develop further readouts involving clock output genes, as the here assessed *Bmal1-luc* and *Per2-luc* reporter profiles, although definitely informative regarding the core clock, might be of limited value as readout for metabolic perturbations.

4.3. Basal IL-6 secretion by human skeletal muscle is regulated by the circadian clock

Importantly, our study demonstrates that human skeletal myotubes, synchronized *in vitro* by forskolin or dexamethasone, secrete basal IL-6 in a circadian manner (Figure 3A,D, Supplementary Figure 5). Furthermore, this circadian pattern is flattened under *siClock*-mediated clockwork disruption, with overall basal IL-6 secretion being strongly downregulated by oscillator perturbation (Figure 3A,D, Table 3). The experimental settings we have developed, combining continuous perfusion and bioluminescence recording, allow for the direct assessment of IL-6 secretion by cultured human myotubes. Studies in healthy adults have suggested that IL-6 levels in the cerebrospinal fluid exhibit a 24 h oscillatory profile, and plasma levels of IL-6 have a

biphasic 12 h component [38,39]. However, it has to be taken into account that plasma levels are originating from different sources of IL-6 and are also influenced by absorption in effector organs.

Our finding that basal IL-6 secretion is regulated locally by the skeletal muscle clock is in line with accumulating evidence that endocrine body rhythms are tightly regulated in humans by the circadian system at several levels (reviewed in [40]). Insulin secretion by beta cells in rodents was suggested to be a subject for clock regulation [11]. Moreover, a number of proinflammatory cytokines exhibit pronounced circadian alterations in the magnitude of their response to an endotoxin challenge at different times of the day [41], among them IL-6. Such circadian gating of the inflammatory response was lost for IL-6 in *REV-ERB α* knockout mice. Moreover, attenuation of *REV-ERB α* levels in human macrophages implied a direct link between *REV-ERB α* and IL-6 secretion [41].

Given a general controversy around the roles of IL-6 level alterations in the etiology of metabolic diseases [42], such downregulation might have a positive or negative impact on skeletal muscle function and body metabolism. A recent *in vivo* study in humans suggested that injection of IL-6 to T2D patients did not affect insulin-stimulated glucose uptake [43], while it was reported to have beneficiary effects on the glucose uptake in young, healthy subjects [44]. Indeed,

Table 3 — Overall basal IL-6 secretion after forskolin synchronization in human myotubes.

	Non-obese (<i>n</i> = 3)	Obese (<i>n</i> = 3)	All donors (<i>n</i> = 6)
	Mean ± SEM	Mean ± SEM	Mean ± SEM
IL-6/48h in <i>siControl</i> samples (pg)	4386.54 ± 411.71	5888.25 ± 1547.86	5137.40 ± 796.43
IL-6/48h in <i>siClock</i> samples (pg)	1424.12 ± 408.83	2395.52 ± 625.60	1909.82 ± 385.20
Inhibition of secretion (%)	69.30 ± 10.61	59.66 ± 7.04	64.48 ± 6.09
<i>Clock</i> mRNA silencing (%)	91.32 ± 2.67	78.57 ± 8.15	84.94 ± 4.57

acute high levels of IL-6 found after exercise improve insulin sensitivity, glucose metabolism [18,45], and fat metabolism [46], whereas chronic exposure to IL-6 causes insulin resistance in mice [47]. At the same time, *IL6* knockout mice develop mature-onset obesity and glucose intolerance [48]. We speculate that similar to cortisol, thyroid hormones, insulin, and other key hormones, which exhibit circadian rhythmicity of their levels [40,49], the oscillatory pattern of IL-6 secretion by the skeletal muscle might give an advantage for the temporal coordination of this myokine with the rest–activity cycle, thus ensuring optimal IL-6 response to external cues.

Our results presented in Figure 3C suggest that the circadian regulation of basal IL-6 secretion may not occur at the transcriptional level. As it has been reported, the clock is quite insensitive to large fluctuations of transcription rate [50]. Moreover, in mouse livers almost 50% of the cycling proteins do not have rhythmic steady-state mRNA levels [51,52]. Similarly, the *IL6* transcriptional profile was not circadian in cultured mHypoE-37 neurons [53], while a low amplitude oscillatory profile of *IL6* transcription was detected in human monocyte-derived macrophages [41] and in mouse astrocytes, where *IL6* transcription was suggested to be directly regulated by ROR α [41,54]. Application of more sensitive transcription analysis approaches (RNA sequencing in samples with a higher temporal resolution), as well as addressing the potential regulation of *IL6* transcription in human skeletal muscle in particular by REV-ERB α and ROR α , might shed additional light on this conjunction. As IL-6 is extensively modified at the post-translational level, one plausible hypothesis beyond transcription could be that the circadian profile of IL-6 secretion is controlled by post-translational modification [55].

Visualizing IL-6 secretion at the single cell level in synchronized human myotubes would be necessary to provide an insight into the mechanism of this phenomenon.

4.4. Functional circadian oscillator in human skeletal myotubes is required for proper basal secretion of a broader range of myokines

Of note, our screening for additional clock-controlled myokines, employing a human cytokine array, suggests that the regulatory role of the skeletal muscle clock is not restricted solely to basal IL-6 secretion but might represent a more general mechanism involved in the fine-tuning of a panel of myokines (Figure 4, Table 4). Specifically, the temporal profiles and/or overall levels of basal secretion of IL-8, MCP-1, M-CSF, GRO- α , VEGF, CD44, Galectin-3, FABP3 and TIMP-1 by human skeletal myotubes imply that these myokines might be regulated by the circadian clock (Figure 4, Table 4).

The following limitations of this part of the study should be taken into account: the low number of around-the-clock perfusion experiments analyzed by multiplex, due to the high experimental complexity of the automated perfusion system, and the low basal concentration levels of many myokines. Therefore, although the technical reproducibility for the duplicates in each around-the-clock experiment was high, in view of the large variability among the human donors with respect to their myokine secretion levels, these experiments must be interpreted with caution. More experimental repetitions will therefore be required to claim the circadian regulation of the myokines identified in our screen. Furthermore, in order to allow for the detection and quantification of myokines with low concentration levels in a circadian manner, more sensitive tools need to be developed.

Table 4 — Myokines with clock-regulated basal secretion.

	Circadian analysis		Fold change of secretion		
	<i>(p</i> -values calculated by JTK_CYCLE, <i>n</i> = 3)		<i>(mean</i> ± SEM, <i>n</i> = 3)		
	<i>siControl</i>	<i>siClock</i>	<i>siControl</i>	<i>siClock</i>	Paired t-test
CD44 ^a	0.777	0.692	1.00	0.67 ± 0.13	0.025*
CHI3L1/YKL40 ^a	1.000	1.000	1.00	2.12 ± 1.22	0.378
FABP3/H-FABP ^a	1.000	0.231	1.00	0.53 ± 0.11	0.002**
Galectin-3 ^a	1.000	0.096	1.00	0.41 ± 0.07	5.19E-6***
GRO- α /CXCL1 ^a	0.096	1.000	1.00	0.55 ± 0.23	0.083
IGFBP-3 ^a	1.000	1.000	1.00	0.70 ± 0.23	0.226
IL-6 ^b	0.01*	1.000	1.00	0.44 ± 0.16	0.006**
IL-8 ^b	0.020*	1.000	1.00	0.47 ± 0.20	0.036*
MCP-1/CCL2 ^a	0.020*	0.096	1.00	0.59 ± 0.27	0.157
M-CSF/GSF1 ^a	0.059	0.777	1.00	0.64 ± 0.16	0.053
MMP-2 ^a	1.000	1.000	1.00	0.78 ± 0.19	0.268
Serpin E1/PAI-1 ^a	1.000	1.000	1.00	0.69 ± 0.16	0.070
Serpin C1 ^a	0.492	1.000	1.00	0.64 ± 0.19	0.123
TIMP-1 ^a	1.000	1.000	1.00	0.68 ± 0.14	0.043*
VEGF ^b	0.949	0.231	1.00	0.60 ± 0.10	0.002**

IFN- γ ^b, IL-1 beta^b, IL-2^b, IL-4^b, IL-7^a, IL-10^b, IL-12p70^b, IL-13^a, IL-17A^a, TNF alpha^b were below detection level.

p* < 0.05, *p* < 0.01, ****p* < 0.001.

^a custom-made luminex screening plate.

^b high sensitivity luminex performance plate.

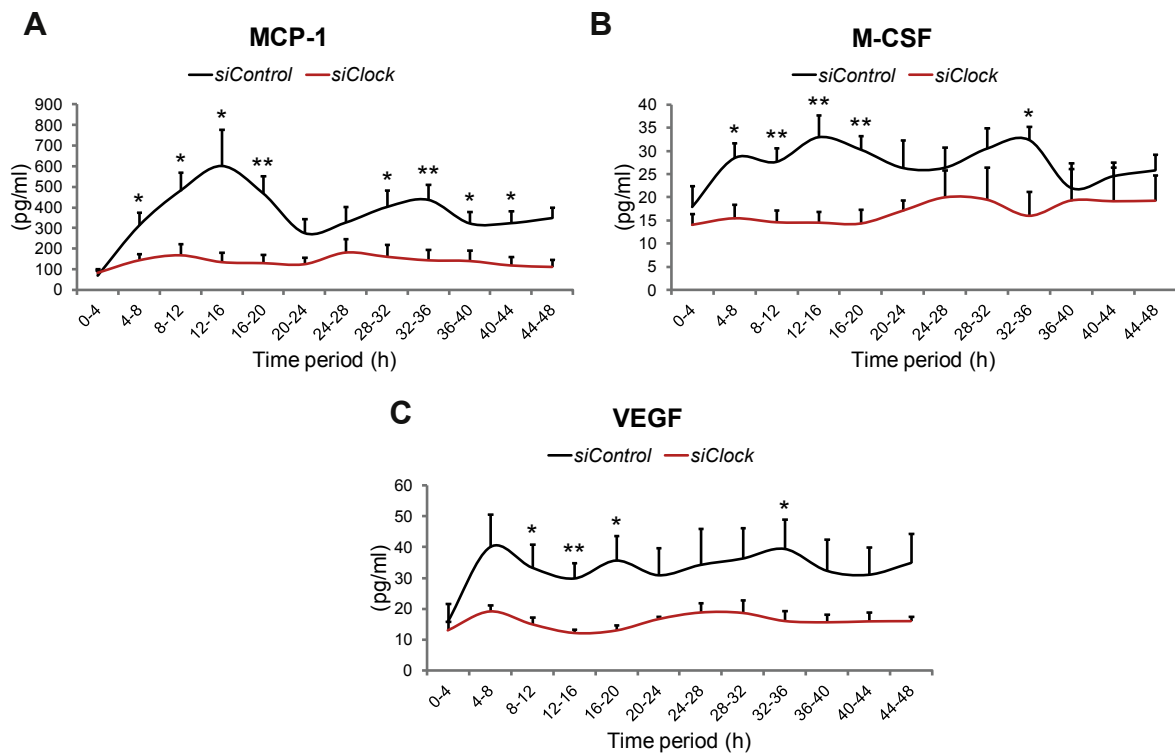


Figure 4: Basal myokine secretion by human skeletal myotubes is affected by circadian clock disruption. Myoblasts were transduced with the *Bmal1-luc* lentivector, differentiated into myotubes, transfected with either *siControl* or *siClock* siRNA, and subjected to continuous perfusion with parallel bioluminescence recording. Concentrated perfusion samples were assessed by multiplex analysis. 2 technical duplicates from 3 biological samples were analyzed for each time point, and normalized to the total DNA content. Basal secretion profiles (mean + SEM) in the presence or absence of a functional clock are shown for (A) MCP-1, (B) M-CSF, and (C) VEGF.

Of note, myokines identified in our multiplex screen as potentially clock-regulated for their basal secretion (Table 4, Figure 4) have been previously linked to obesity or T2D. For instance, serum IL-8 levels are increased in T2D patients, with a significant diurnal variation observed for IL-8 in blood upon LPS-stimulation [56,57]. Furthermore, insulin-resistant human myotubes secrete higher levels of IL-8 [19]. MCP-1, which mediates skeletal muscle macrophage recruitment, was also suggested to play a role in the etiology of T2D [20]. Interestingly, this proinflammatory cytokine has been previously shown to exhibit pronounced circadian alterations in the magnitude of its response to endotoxin challenge at different times of the day [41]. M-CSF has been shown to play an important role in inflammatory diseases including obesity [58]. VEGF, known for its angiogenic properties, is transcriptionally regulated by the circadian clock [59], and has also been implicated in T2D and insulin resistance [60,61]. Plasma levels of CD44 were positively correlated with insulin resistance in humans [62,63]. Furthermore, genetic association studies have linked CD44 with T2D [62,64]. Taken together, our data suggest that disruption of the circadian clock might affect the level and temporal profile of basal secretion for a number of myokines that play a role in the etiology of T2D and obesity. Inevitably, our findings raise the question on the physiological relevance of basal myokine secretion. One plausible argument is that while induced myokine secretion is regulated acutely (for instance by exercise), circadian regulation of basal myokine secretion might represent a fine-tuning mechanism, allowing adaptation of the skeletal muscle function to the rest–activity cycle. Given the major role of the circadian clock in allowing organisms to anticipate daily environmental

changes rather than react to them, circadian regulation of basal myokine secretion might represent such an anticipatory mechanism that coordinates skeletal muscle “availability”. Furthermore, this newly discovered link between the functional skeletal muscle clock and basal secretion of a number of myokines might bear potential consequences for the development of chronic diseases, such as obesity and T2D.

5. CONCLUSION

Skeletal muscle represents the most important site of insulin resistance in T2D patients [12]. Moreover, the emerging critical role of inflammation in the etiology of T2D makes inflammatory cytokines plausible candidates for developing new therapeutic approaches for the treatment of this disease [65]. It is therefore of highest scientific and clinical importance to provide further insight into the emerging connection between circadian oscillator function, metabolic regulation, and T2D. Given that human primary myotubes, established from T2D patient biopsies and cultured *in vitro*, have been demonstrated to maintain their *in vivo* phenotypes of inflammation and insulin resistance [66], our model of synchronized cultured primary myotubes represents a valuable experimental tool that allows for studying the role of the circadian oscillator in human skeletal muscle function upon metabolic diseases. This work is the first detailed characterization of the human skeletal myotube circadian oscillator and its critical impact on basal myokine secretion. It opens the way for future studies that may link defects in these pathways with insulin resistance, obesity, and T2D. Given obvious obstacles for studying the human circadian

oscillator *in vivo*, our experimental approach, using human primary cell cultures established from biopsies that express a luciferase reporter driven from a circadian promoter, constitutes a powerful model system for human skeletal muscle clock studies.

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CONFLICT OF INTEREST

The authors have no conflict of interest to declare.

APPENDIX A. SUPPLEMENTARY DATA

Supplementary data related to this article can be found at <http://dx.doi.org/10.1016/j.molmet.2015.07.009>.

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