The circadian clock regulates rhythmic activation of the NRF2/glutathione-mediated antioxidant defense pathway to modulate pulmonary fibrosis

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The disruption of the NRF2 (nuclear factor erythroid-derived 2-like 2)/glutathione-mediated antioxidant defense pathway is a critical step in the pathogenesis of several chronic pulmonary diseases and cancer. While the mechanism of NRF2 activation upon oxidative stress has been widely investigated, little is known about the endogenous signals that regulate the NRF2 pathway in lung physiology and pathology. Here we show that an E-box-mediated circadian rhythm of NRF2 protein is essential in regulating the rhythmic expression of antioxidant genes involved in glutathione redox homeostasis in the mouse lung. Using an in vivo bleomycin-induced lung fibrosis model, we reveal a clock “gated” pulmonary response to oxidative injury, with a more severe fibrotic effect when bleomycin was applied at a circadian nadir in NRF2 levels. Timed administration of sulforaphane, an NRF2 activator, significantly blocked this phenotype. Moreover, in the lungs of the arrhythmic Clock<sup>D19</sup> mice, the levels of NRF2 and the reduced glutathione are constitutively low, associated with increased protein oxidative damage and a spontaneous fibrotic-like pulmonary phenotype. Our findings reveal a pivotal role for the circadian control of the NRF2/glutathione pathway in combating oxidative/fibrotic lung damage, which might prompt new chronotherapeutic strategies for the treatment of human lung diseases, including idiopathic pulmonary fibrosis.

[Keywords: circadian clock, glutathione, NRF2, bleomycin, pulmonary fibrosis]

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inevitably expose the mammalian cells and tissues to periodic challenges, including oxidative insults from environmental toxins/pollutants and endogenously produced reactive metabolites as products of respiration (Patel et al. 2014). Failure to scavenge reactive oxygen species (ROS), reactive nitrogen species (RNS), and reactive electrophiles results in excessive oxidative stress, leading to damage to critical macromolecules and cellular structures (Veal et al. 2007). The ability to anticipate and withstand such cyclic insults (e.g., through effective ROS scavenging) is therefore essential for normal protective tissue functions. There is emerging evidence that circadian clocks regulate processes that keep ROS at physiological levels and protect organisms from oxidative stress (Stangherlin and Reddy 2013; Patel et al. 2014). Indeed, self-sustained rhythms in the cellular redox state (Stangherlin and Reddy 2013; Patel et al. 2014). The current paradigm of NRF2 regulation has mainly focused on oxidative stress-induced post-translational mechanisms, including regulation by its negative regulators (KEAP1, BACH1, and β-TrCP), positive regulators [DJ1, p62, and p21], or protein modifications [Itoh et al. 1999; McMahon et al. 2003; Chen et al. 2009; Komatsu et al. 2010; Chowdhry et al. 2013]. In response to oxidative stress, NRF2 protein translocates to the nucleus and binds to the antioxidant response elements (AREs) in the promoters of many antioxidant genes. Subsequently, NRF2 induces a transcriptional program that maintains cellular redox balance and protects cells from oxidative insults (Reddy et al. 2007; Malhotra et al. 2010). NRF2 transcriptional targets include glutathione cysteine ligase (GCL), involved in the rate-limiting step of biosynthesis of antioxidant glutathione; glutathione-utilizing enzymes such as glutathione S-transferase (GST); and haeme oxygenase 1 (HMOX1), involved in the catabolism of pro-oxidant haeme.

Nrf2-deficient mice show a decrease in both constitutive and inducible expression of multiple glutathione-dependent enzymes [McMahon et al. 2001], associated with an impaired antioxidant defense and susceptibility to stress-induced tissue pathologies (Cho et al. 2006; Chan and Kan 1999). Consequently, they are highly sensitive to many environmental toxicants and chemical inducers of oxidant injury and fibrosis in several tissues, including bleomycin-induced and butylated hydroxytoluene (BHT)-induced pulmonary fibrosis (Chan and Kan 1999; Cho et al. 2004; Kikuchi et al. 2010). In humans, altered NRF2 expression and impaired redox balance have been associated with the pathogenesis of chronic lung diseases (asthma, COPD, and idiopathic pulmonary fibrosis) as well as lung cancer (Cho et al. 2006; Hayes and McMahon 2009). Indeed, pharmacological targeting of NRF2 has emerged as a novel therapeutic approach to combat oxidative stress seen in chronic lung diseases (Malhotra et al. 2011; Artaud-Macari et al. 2013).

Recently, mRNA levels of Nrf2 as well as several of its target genes have been reported to show diurnal variation in mouse livers and Drosophila heads [Beaver et al. 2012; Xu et al. 2012]. However, whether and how endogenous circadian signals control NRF2-dependent antioxidant activity in tissue physiology and upon pathological challenges, especially in the lung, has not been defined. Here we investigated the circadian rhythms in the protein levels of NRF2 and activity of the NRF2/GSH pathway in lung tissues from light/dark cycle-entrained mice. We also report the physiological and pathological importance for the circadian control of the NRF2/GSH pathway using a well-established lung fibrosis model following a bleomycin challenge. Our data reveal a pivotal role for endogenous NRF2 rhythms in the lung in coupling protective antioxidant responses to the time-of-day susceptibility to oxidant injury and pulmonary fibrosis.

Results

Circadian protein expression of NRF2 transcription factor in the mouse lung

To investigate the hypothesis that the NRF2-mediated antioxidant pathway is under circadian regulation in the lung, we first examined the total NRF2 protein levels in mouse lungs harvested at 4-h intervals from mice kept in constant darkness (DD). Western blotting using a specific NRF2 antibody [confirmed in Nrf2 knockout mouse embryonic fibroblasts (MEFs)] [data not shown] revealed a robust rhythmic pattern of NRF2 protein [one-way ANOVA, Tukey test, P < 0.05]. The NRF2 protein peaked at the circadian time 3–7 (CT3–CT7), reaching a trough at CT15–CT19 (Fig. 1A). Nuclear accumulation of NRF2 protein is essential for its subsequent activation of downstream genes [Reddy et al. 2007; Nguyen et al. 2009]. Indeed, we detected rhythmic levels of NRF2 protein in the nuclear lung fractions, with the maximal levels corresponding to the peak of the total pool of NRF2 protein [P < 0.05] [Fig. 1B]. Immunostaining of NRF2 revealed a significant time-of-day difference in the lungs that was mainly localized to the bronchial and alveolar epithelium, with an approximately threefold weaker signal at “zeitgeber” time 12 (ZT12; lights off) compared with ZT0 (lights on) [Fig. 1C]. The rhythmic NRF2 protein expression was also evident in the total cell lysates [Fig. 1D] and nuclei [Fig. 1E,F] of clock-synchronized Rat1 fibroblasts, demonstrating the cell-autonomous nature of the NRF2 protein rhythms.

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The circadian clock exerts transcriptional control of the Nrf2 gene via a putative E-box element in its promoter

The circadian clock transcription factors regulate target genes through the cis-acting elements in their gene promoters (Panda et al. 2002; Rey et al. 2011). We therefore analyzed the proximal promoter regions of the mouse, rat, and human Nrf2 genes for clock-regulated transcriptional elements, which revealed a conserved putative E-box (CACGTG) close to their transcriptional start sites (TSSs; -704 base pairs [bp] relative to the TSS in the mouse Nrf2) (Fig. 2A). Next, we cloned the wild-type mouse Nrf2::luc promoter (1.7 kb) and generated an E-box mutated construct. Coexpression of circadian transcription activators BMAL1 and CLOCK in Rat1 cells led to a significant stimulation of Nrf2::luc activity (2.1-fold, $P < 0.05$), similar to the activation of the Per1::luc promoter [containing three canonical E-box elements; 2.4-fold, $P < 0.01$] (Fig. 2B). Mutation of the core E-box sequence in the Nrf2 promoter completely abolished its induction by CLOCK/BMAL1 complexes (Fig. 2C). Moreover, coexpression of circadian transcription repressors, either PER2 or CRY1, suppressed the CLOCK/BMAL1 transcriptional activation (Fig. 2D). Finally, stable transfection of the Nrf2::luc reporter into Rat1 cells revealed a clearly discernible circadian rhythm with a period of 23.7 h (data not shown).

We next examined the temporal profiles of endogenous Nrf2 mRNA in the lungs of wild-type and Clock$^{-/-}$ mice [a model in which the positive arm of the clock is genetically disrupted] (Vitaterna et al. 1994). This revealed a clear rhythmic expression of Nrf2 transcripts in wild-type lungs ($P < 0.05$, one way ANOVA) (Fig. 2E). In
contrast, the mRNA rhythm of Nrf2 was abolished and remained low across all time points in the lungs of Clock\textsuperscript{D19} mice, consistent with a loss of rhythms in the clock gene expression (Fig. 2E; Supplemental Fig. S1). Consistently, knockdown of Bmal1 by siRNA significantly reduced the levels of Nrf2 mRNA in Rat1 fibroblasts \( P < 0.05 \) (Supplemental Fig. S2). In contrast, in synchronized MEFs from arrhythmic Cry\textsuperscript{1−/−}/Cry\textsuperscript{2−/−} mice, a model in which the negative arm of the clock is disrupted (Supplemental Fig. S3A,B; van der Horst et al. 1999), we observed constitutively high protein expression levels of NRF2 as well as mRNA levels of its two target genes \( Gclm \) and \( Gsta3 \) (Supplemental Fig. S3C,D).

To verify the function of the putative E-box element in the Nrf2 promoter in vivo, temporal chromatin immunoprecipitation (ChIP) assays were performed in the wild-type and Clock\textsuperscript{D19} lungs. Data [mean ± SEM] were normalized to Gapdh. \( (*) P < 0.05 \), \( (**) P < 0.01 \), t-test. \( (\text{ns}) \) not significant for Clock\textsuperscript{D19} lung, one-way ANOVA for the effect of time. (F) Temporal CLOCK binding to E-boxes in Nrf2 and Dbp gene promoters in wild-type and Clock\textsuperscript{D19} lungs using CLOCK-specific ChIP. IgG served as a negative control. Data [mean ± SEM] were expressed as percent input. \( (*) P < 0.05 \), \( (\text{ns}) \) not significant, t-test. (White bar) Light phase; (black bar) dark phase.
pancy was also observed in the E-box region of the Nrf2 promoter (Fig. 2F) but not in the distal region of the promoter (Supplemental Fig. S4A). There was a significantly higher binding of CLOCK at ZT4 as compared with ZT16, coinciding with the peak and trough expression of Nrf2 mRNA, respectively. Moreover, CLOCK binding to the Nrf2 promoter was arrhythmic and remained at an intermediate level in the Clock−/− lungs (Fig. 2F). Together, these results demonstrate that the Nrf2 gene is directly regulated by the core clock components in vitro and in vivo through the conserved E-box element in its promoter.

**Rhythmic promoter binding of NRF2 drives oscillations of ARE-regulated antioxidant genes involved in glutathione homeostasis**

Upon nuclear translocation, NRF2 protein binds to the AREs in the promoters of its target genes and thus maintains both basal and inducible expression of many antioxidant genes (Nguyen et al. 2009; Malhotra et al. 2010). The rhythmic nuclear NRF2 levels prompted us to examine whether the expression of ARE-regulated NRF2 target genes is also rhythmic. Temporal quantitative PCR (qPCR) of the mouse lungs demonstrated robust endogenous circadian rhythms for a number of Nrf2 target genes, including Gclm, Gsta3, and Hmox1 (P < 0.05) [Fig. 3A; Supplemental Fig. S5], as well as a core clock gene, Bmal1 (P < 0.001) (Supplemental Fig. S5). To investigate the hypothesis that NRF2 may be rhythmically recruited to the AREs in the promoters of antioxidant genes in vivo, we performed temporal ChIP assays in the mouse lungs using an anti-NRF2 antibody. Here, we detected strongly rhythmic NRF2 binding specifically to the promoter regions spanning the well-characterized AREs [Malhotra et al. 2010] in Gclm and Gsta3 gene promoters (Fig. 3B; Supplemental Fig. S4B). We observed significantly higher promoter occupancy at ZT0 and ZT16 [light phase] and lower occupancy at ZT12 and ZT18 [dark phase]. The peak of NRF2 binding during light phase is in agreement with its maximal nuclear expression and the peaks of Gclm and Hmox1 mRNAs in the lung. Interestingly, despite maximal NRF2 binding to the Gsta3 promoter during the light phase, Gsta3 mRNA peaked during early subjective night, suggesting that additional phase delay mechanisms may be at play, such as the availability of

**Figure 3.** Circadian rhythm of NRF2 binding to ARE drives rhythmic oscillations of genes involved in glutathione synthesis and utilization. (A) Temporal mRNA levels of NRF2 targets in wild-type [WT] mouse lungs [DD]. Data [mean ± SEM] were normalized to Gapdh, and the lowest expression was set as 1. (**) P < 0.05, one-way ANOVA. (B) Rhythmic NRF2 occupancy on AREs of antioxidant gene promoters in wild-type mouse lungs by NRF2-specific ChIP. The position of each ARE in relation to the TSS is shown. Data [mean ± SEM] were normalized to Gapdh. (**) P < 0.05 for all six genes in wild type; not significant for Nrf2 knockout MEFs (except Hmox1 and Dpb; [*] P < 0.05), one-way ANOVA for the effect of time. (C) Temporal levels of reduced glutathione (GSH) in MEFs from wild-type and Nrf2 knockout mice following serum shock. Data [mean ± SEM] were normalized to cellular counts and quantified using standard curve for reduced GSH. (***) P < 0.01; [ns] not significant, t-test.
cofactors, as shown for the transcription factor BMAL1 (Rey et al. 2011), or post-transcriptional processing (Patel et al. 2014). The difference in the timing of maximal expression may be necessary for the specific function of individual NRF2 targets.

To investigate whether NRF2 is required for the rhythmic expression of antioxidant genes, we performed temporal qPCR for a panel of known NRF2 target genes in clock-synchronized MEFs from wild-type and Nrf2 knockout mice. Here, we observed significantly reduced basal expression of many antioxidant genes in Nrf2 knockout MEFs, as reported previously (Malhotra et al. 2010). Interestingly, the rhythmic patterns of genes coding for enzymes involved in glutathione synthesis (Gclm and Gclc) and glutathione utilization (Gsta3 and Gsta4) were either severely dampened or abolished (Fig. 3C). In contrast, Hmox1, an inducible (but not basal) NRF2 target, remained robustly rhythmic in Nrf2 knockout MEFs. Consistent with these results, the level of total glutathione (GSH) showed a significant temporal variation in wild-type MEFs but remained constitutively low in Nrf2 knockout MEFs [Fig. 3D]. Importantly, the rhythmic expression of Ddbp in Nrf2 knockout MEFs [Fig. 3C] and the behavioral free-running rhythms appear preserved in Nrf2 knockout mice (Supplemental Fig. S6). These findings support a direct NRF2-dependent rhythmic control of its downstream targets rather than due to feedback actions of NRF2 on the molecular circadian clock. All together, these data demonstrate that the periodic NRF2 activity regulates the rhythmic GSH homeostasis through temporally controlling a key set of genes involved in glutathione synthesis and utilization.

Time-of-day-dependent lung response to bleomycin challenge is coupled to temporal NRF2 activity

The NRF2 antioxidant pathway is known to play a critical role in combating fibrotic injury (Cho et al. 2004; Walters and Kleeberger 2008). We therefore hypothesized that the rhythmic NRF2 activity may gate the tissue-protective responses to a fibrotic challenge in a time-of-day-dependent manner. To test this, we used an established in vivo mouse model of bleomycin-induced lung fibrosis (Walters and Kleeberger 2008). Bleomycin was administrated intracheally close to either the peak or nadir (ZT0 or ZT12, respectively) of NRF2 protein expression [see Fig. 1A–C]. As expected, 7 d after bleomycin challenge, characteristic histological changes, including areas of inflammatory cell infiltration and alveolar wall thickening [data not shown], were observed at both time points. In addition, we also observed a significant increase in collagen accumulation around the peri-bronchial regions, as revealed by Masson’s Trichrome staining of lung sections and quantification of fibrosis using the Ashcroft scoring system [Fig. 4A; Ashcroft et al. 1988]. However, the degree of lung fibrosis was clearly dependent on the time of treatment and inversely correlated with NRF2 protein levels. A significantly higher fibrotic score was observed when bleomycin was delivered at ZT12 as compared with ZT0 [Fig. 4B], associated with higher induction of fibrotic gene markers (Timp1, Col1a2, and Mmp3) [Supplemental Fig. S7]. Bleomycin challenge is known to activate NRF2-dependent antioxidant tissue responses (Cho et al. 2004). Following an acute bleomycin challenge (4 h), the induction of mRNAs for Gclc, Gsta3, and Hmox1 showed a clear time-of-day dependence, with a significantly higher induction at ZT0 than at ZT12 [P < 0.01 for all three genes] [Fig. 4C], consistent with the hypothesis of a clock “gated” antioxidant tissue response.

To further establish the role of NRF2 rhythms in time-of-day susceptibility to bleomycin-induced fibrosis, we investigated whether the timed administration of sulforaphane (SFN; a direct NRF2 activator) can reduce the degree of bleomycin-induced fibrosis seen at the circadian nadir in NRF2 levels. To this end, wild-type mice were pretreated with either vehicle or SFN administered intraperitoneally at ZT6 [and thereafter at the same time point on days 1, 3, and 5] prior to a single challenge with bleomycin at ZT12 and assessed 7 d later. As a control, mice treated with SFN alone [at ZT6] showed increased NRF2 protein levels in the lungs at ZT12 as compared with vehicle treatment [data not shown]. Here, the degree of bleomycin-induced lung fibrosis was significantly reduced by the timed SFN treatments [P < 0.05] [Fig. 4D]. Consistent with these findings, the bleomycin-induced expression of fibrotic markers (Col1a2, Timp1, Mmp3, and Ctgf) was significantly blunted by SFN treatment [Fig. 4E]. All together, these results strongly support a role for NRF2 rhythms in coupling antioxidant response to time-of-day susceptibility to bleomycin-induced lung fibrosis. Moreover, these findings suggest that timed SFN treatment might protect the lungs from oxidative challenges, especially those that fall within the nadir of NRF2 activity.

Altered NRF2/GSH pathway activity in the Clock los-19 mouse model to investigate the requirement of the functional circadian clock, we used the Clock los-19 mouse model to investigate changes in the NRF2 pathway activity. Consistent with the loss of Nrf2 mRNA rhythmicity in the Clock los-19 lungs, NRF2 protein expression was constitutively low and arrhythmic [Fig. 5A]. Moreover, ChIP assays revealed an arrhythmic and low NRF2 occupancy in the ARE region of the promoters for Gclm and Gsta3 [Fig. 5B]. While qPCR revealed robust time-of-day-dependent mRNA expression of Gclm and Gsta3 in wild-type lungs, their expression in the Clock los-19 lungs remained constitutively lower, at levels similar to or below the trough expression in wild-type lungs [Fig. 5C].

In an attempt to test the hypothesis that the Clock los-19 lungs were more prone to bleomycin-induced fibrotic injury, unexpectedly, we identified a spontaneous fibrotic-like phenotype in Clock los-19 lungs even without a bleomycin challenge. We found a significantly increased deposition of collagen fibers around the bronchioles in Clock los-19 com-
pared with wild-type lungs \( P < 0.001 \) [Fig. 6A]. Consistently, the expression of fibrotic marker genes \( \text{Mmp3} \) and \( \text{Ctgf} \) was significantly elevated in \( \text{Clock}\textsuperscript{D19} \) lungs [Fig. 6B]. A reduced GSH level is known to be involved in the fibrotic injury and remodeling processes (Liu and Gaston Pravia 2009). To explore the underlying mechanisms for the observed lung phenotype, we compared the temporal levels of reduced glutathione, a major antioxidant output of the NRF2 pathway. In wild-type lungs, we found a significantly higher level of reduced GSH at ZT0 compared with ZT12 (Fig. 6C). In contrast, \( \text{Clock}\textsuperscript{D19} \) lungs showed constitutively lower levels of reduced GSH, equivalent to the nadir level in wild-type lungs. Similar alterations in GSH level were also found in synchronized \( \text{Clock}\textsuperscript{D19} \) MEFs, which could partially be restored by SFN treatment (Supplemental Fig. S8; Higgins et al. 2009), supporting the involvement of NRF2.

To determine whether the low GSH levels in \( \text{Clock}\textsuperscript{D19} \) lungs correlated with an increased oxidative burden, we temporally examined the lung protein carbonylation levels (a marker of oxidative damage). We observed a time-dependent change in the protein carbonyl levels in wild-type lungs, which were anti-phasic to the reduced GSH levels (Fig. 6D). In contrast, protein carbonylation levels remained high at both time points in \( \text{Clock}\textsuperscript{D19} \) lungs, consistent with a constitutive decline in GSH levels. These data support the hypothesis that the functional clock is required in coordinating the temporal activation of the NRF2/GSH pathway in the lung, disruption of which contributes to the increased susceptibility of the lung to cyclic oxidative stress and profibrotic challenges.

**Discussion**

It has recently been suggested that circadian clocks may mediate the protective responses to oxidative stress...
However, the underlying clock-controlled molecular mechanisms and their role in tissue-specific physiology are still poorly understood. In this study, we identified the circadian clock as an endogenous molecular regulator of the NRF2/glutathione-mediated antioxidant defense pathway in the mouse lung, which couples the protective antioxidant response to the time-of-day variation in susceptibility to oxidant injury and pulmonary fibrosis. Based on our findings, we propose a working model depicting how the circadian clock exerts its regulation of protective antioxidant responses in the lung through the control of transcription factor NRF2 under physiological conditions (Fig. 7). The circadian transcription factors CLOCK and BMAL1 positively regulate $Nrf2$ transcription through temporal control of the E-box element in the $Nrf2$ gene promoter. NRF2 protein thus accumulates in a circadian manner and drives circadian transcription of a key set of antioxidant genes involved in glutathione metabolism (such as $Gclm$ and $Gsta3$). The periodic $Nrf2$ activity is exerted through its rhythmic recruitment to the AREs in the promoters of target antioxidant genes and mediates a time-coordinated protection against oxidative tissue injury and fibrotic damage.

Indeed, using an in vivo bleomycin-induced pulmonary fibrosis model, we show for the first time that, in the lung, the acute induction of the NRF2-mediated antioxidant response is temporally “gated” by the clock and coupled to a time-dependent resistance to fibrotic injury. Importantly, a scheduled pharmacological activation of NRF2 in vivo was able to significantly mitigate the decrease in resistance to fibrotic effects of bleomycin delivered at the circadian nadir in NRF2 levels. Moreover, in the lungs of $Clock^{Δ19}$ circadian mutant mice, the altered temporal regulation of NRF2 antioxidant activity was associated with diminished glutathione levels, increased oxidative protein damage, and a spontaneous fibrotic-like phenotype. All together, our findings highlight an important role of the circadian clock in the regulation of the NRF2/GSH-mediated antioxidant response and its coupling to oxidative/fibrotic injury in the lung.

Our study builds on the recent reports implicating the role of the circadian clock in the regulation of tissue physiology through redox-dependent mechanisms [Kondratov et al. 2006; Lai et al. 2012; Wang et al. 2012; Musiek et al. 2013; Peek et al. 2013]. For instance, direct circadian regulation of antioxidant genes [$Ngo1$ and $Aldh2$] has been reported in the cerebral cortex, and the circadian regulation of NAD$^+$-dependent histone deacetylase SIRT3 has been implicated in mitochondrial oxidative metabolism. Our findings reveal a novel aspect of circadian clock function in modulating pulmonary fibrotic responses, which is mediated through its control of the NRF2 transcription factor.

Current research concerning NRF2 regulation has largely focused on the post-translational mechanisms, especially via a KEAP1-based protein turnover in response to various electrophilic/oxidative stresses. In this model, NRF2 proteins are tethered to the cytoplasm by...
KEAP1 and targeted for proteasomal degradation (Hayes and McMahon 2009). It was therefore thought that only a small pool of NRF2 protein can escape from its degradation in the cytosol and translocate to the nucleus under basal conditions. In response to oxidative stress, however, oxidative inactivation of KEAP1 and de novo synthesis of NRF2 drive sustained nuclear NRF2 accumulation (Itoh et al. 2010). Our data reveal a rhythmic nuclear abundance of endogenous NRF2 protein in tissues and cells that accompanies the rhythmic levels in the total pool of NRF2 protein. These results are supported by previous studies in the mouse liver and pancreatic β cells that identified Nrf2 as one of the clock-controlled targets (Xu et al. 2012; Lee et al. 2013). It is therefore conceivable that during the circadian peak of NRF2 protein, increased levels of NRF2 may saturate the binding capacity of KEAP1, thus allowing NRF2 to escape from the KEAP1-mediated degradation and translocate to the nucleus, where it induces higher expression of antioxidant genes. In contrast, during the nadir phase of NRF2 expression, KEAP1 may exert tighter control of NRF2 activity through increased NRF2 degradation. In this way, the regulation of NRF2 nuclear activity appears to be restricted to a particular circadian window, gated by both the clock-controlled mRNA synthesis and the KEAP1-based protein degradation.

Previous ChIP-seq (ChIP combined with deep sequencing) and microarray studies have revealed that, in addition to stress-induced antioxidant gene expression, NRF2 directly controls basal expression of numerous antioxidant genes via AREs in their promoters (Reddy et al. 2007; Malhotra et al. 2010). Here we show that, under physiological conditions in the lung, the protein levels and recruitment of NRF2 to gene-specific antioxidant gene promoters is under rhythmic clock regulation. Nrf2-deficient MEFs are not capable of maintaining circadian rhythms of a number of key antioxidant genes involved in glutathione homeostasis and show arrhythmic and reduced levels of its major antioxidant output glutathi-
one. Moreover, in ClockΔ19 circadian mutant mice, the altered regulation of NRF2 activity was also associated with reduced expression of genes involved in glutathione metabolism as well as total glutathione levels. Since both the behavioral free-running rhythms in Nrf2 knockout mice and Dhp mRNA rhythm in Nrf2 knockout MEFS appear preserved, this highlights the essential role of NRF2 in mediating rhythmic glutathione homeostasis.

Changes in the lung NRF2 expression and GSH metabolism as well as elevated protein carbonyl content are recognized as central features of many human chronic lung diseases, including idiopathic pulmonary fibrosis, cystic fibrosis, acute respiratory distress syndrome, and COPD (Rahman and MacNee 2000; Dalle-Donne et al. Rangasamy et al. 2004; Reddy et al. 2007; Liu and Gaston Pravia 2009). NRF2 activity was associated with constitutively lower one metabolism as well as total glutathione levels. Since both the behavioral free-running rhythms in Nrf2 knockout mice and Dhp mRNA rhythm in Nrf2 knockout MEFS appear preserved, this highlights the essential role of NRF2 in mediating rhythmic glutathione homeostasis.

In conclusion, we identified the molecular circadian clock as an endogenous regulatory mechanism controlling the rhythmic activity of the redox-sensitive transcription factor NRF2 in the mouse lung. We also reveal the physiological significance of this rhythmic control in modulating tissue susceptibility to oxidative injury and pulmonary fibrosis. Given the role of NRF2 in human fibrotic diseases, we envisage a scenario in which circadian misalignments [e.g., caused by genetic variation or aging] may compromise the rhythmic activation of the NRF2-mediated antioxidant defense, leading to increased oxidative damage. Moreover, once fibrosis develops, there is no way to revert the phenotype. Current therapy that aims to slow down the progression of fibrotic diseases is often ineffective (Liu and Gaston Pravia 2009); our study implicates the need for considering circadian timing mechanisms, including timed drug administration (chronopharmacology) (Levi and Schibler 2007). Therefore, a major challenge in the future will be to understand how these circadian mechanisms affect tissue function in human disease.

Materials and methods

Animals and tissue collection

All experiments were conducted under the aegis of the 1986 Home Office Animal Procedures Act [UK]. Mice were maintained on a standard maintenance chow under a 12-h light/12-h dark (12:12 LD) regimen. PER2::Luc and ClockΔ19 mice on a C57BL/6J background were generated by Professor Joseph Takahashi. ClockΔ19 mice were subsequently bred with PER2::Luc mice. For circadian tissue collections or experiments with bleomycin-induced fibrosis, 8-wk-old female C57BL/6J mice (Harlan Laboratories) were placed under 12:12 LD cycles for 2 wk before their release into DD. Animals were sacrificed by cervical dislocation in complete darkness using an infrared viewer, and lung tissues were harvested at 4-h intervals, beginning at 39 h after the start of DD. All tissues [circadian and diurnal collections] were either freshly used or snap-frozen in liquid nitrogen and kept at −80°C until use.

Reagents and plasmids

Bleomycin was purchased from ENZO. SFN was purchased from LKT Laboratories. All other chemicals were purchased from Sigma unless specified otherwise. mCLOCK and hBMAL1 expression plasmids and pGL3-mPer1::luc were kind gifts from Dr. Kazuhiro Yagita. The lentiviral mPer2::luc construct was previously described (Lu et al. 2010). The generation of wild-type and ΔE-box mNrf2::luc constructs is described in the Supplemental Material.

Real-time bioluminescence recordings

Bioluminescence from the organotypic lung slices of wild-type and ClockΔ19 mice [on a PER2::Luc background] [Yoo et al. 2004] or from MEFS with lentivirally transduced Per2::luc
reporters was recorded by photomultiplier tubes (PMTs; Hamamatsu) housed in a light-tight incubator at 37°C and 0% CO2 as described previously [Meng et al. 2008]. The lung tissue slices were cultured in the recording medium [phenol red-free DMEM supplemented with 10% fetal bovine serum, 3.5 mg/mL glucose, 25 U/mL penicillin, 25 μg/mL streptomycin, 0.1 mM luciferin, 10 mM HEPES-NaOH at pH 7] on Millipore cell culture inserts [Millipore]. Cellular clocks were synchronized by a well-established serum shock protocol using 50% horse serum [for 2 h]. Data were presented as photons counts per minute.

ChIP assays

Freshly harvested lung tissues were homogenized and cross-linked in 1% [v/v] formaldehyde. Lung nuclear lysates were sonicated (~500 bp) using Bioruptor (Diagenode), and 5% of each sheared chromatin sample was saved as an input control. Following preclearing, lung chromatin was incubated overnight at 4°C with the following antibodies: 1 μg/mL rabbit anti-NRF2 (Santa Cruz Biotechnology), 4 μg/mL mouse anti-CLOCK (CLSP3) (Yoshitane et al. 2009, in-house), and 1 μg/mL control rabbit or mouse IgGs [Millipore]. Immunoprecipitated DNA fragments were captured using protein G magnetic Dynabeads [Life Technologies]. After a series of washes, proteinase K digestion, and reverse cross-linking, DNA was eluted and cleaned up (Qiagen). Real-time qPCR was performed using TaqMan-based primer/pairs [Supplemental Table 1] to amplify the promoter regions spanning the conserved E-box or AREs. The amount of precipitated DNA in each sample was quantified according to the percent input method, in which Ct values of each sample were normalized to the adjusted Ct values of their respective inputs (corrected for dilution factor) and expressed as percent input.

Tissue glutathione (GSH) assay

Frozen lung tissues [100 mg] were homogenized in ice-cold 5% sulfoasacilic acid [Sigma] to minimize oxidation of reduced GSH. The diluted acid supernatants were used for GSH measurements, and the acid-precipitated protein was used for protein determination using a BCA protein assay kit [Pierce]. Free reduced glutathione was determined using the luminescence-based GSH-Glow glutathione assay [Promega], which is based on the conversion of a luminogenic probe in the presence of reduced GSH in a reaction catalyzed by GST. Luminescence measurements were performed in triplicates for each biological replicate and processed in a single run to aid comparative quantification. Relative luminescence units (RLUs) were corrected for background luminescence and converted to nanomoles per milligram of protein using a standard curve for reduced GSH.

Protein carbonylation assay

To measure protein carbonylation, lung protein lysates were processed using the Oxyblot protein oxidation detection kit [Millipore]. Briefly, the carbonyl groups in the SDS-denatured proteins were derivatized to 2,3-dinitrophenyl hydrazone (DNP-hydrazone) by reaction with 2,4-dinitrophenylhydrazine [DNPH] followed by SDS-PAGE. Immunoblot detection of protein carbonyls was performed using rabbit anti-DNP primary antibody and goat anti-rabbit HRP-conjugated secondary antibody. As a negative control, proteins were derivatized using a derivatization control solution. Protein carbonylation levels were analyzed using National Institutes of Health ImageJ software by determining the values obtained for total protein density within each protein lane and were normalized to tubulin density for each sample.

Bleomycin-induced lung fibrosis model

All animal procedures were carried out in accordance with the Animals Scientific Procedures Act (1986). Female wild-type (C57/BL6), PER2::Luc, or clock-tau19 mice [aged 8–12 wk] were dosed intratracheally with 1 mg/kg clinical-grade bleomycin [30 μL volume, ENZO] or vehicle [saline] at either ZT0 [lights on] or ZT12 [lights off]. Tissues were collected either 4 h later for RNA analysis of acute gene transcription or 7 d later [at ZT6] for histological assessment and long-term gene expression. The lungs were lavaged post-mortem via an incision in the trachea, with 1 mL of 10 mM EDTA and 1% BSA. For acute gene expression assays, fresh lung lobes were dissected and snap-frozen. For histological samples, 1 mL of 4% PFA was delivered via the trachea to inflate the lungs. The trachea was clamped off, and the lungs were removed en bloc and fixed in 4% PFA overnight at 4°C. After fixating, the middle right lobe was removed for RNA extraction, while the left lobe was processed for histological assessment. Lung slices at 5 μm were stained with Masson’s Trichrome solution following the standard procedure within our core histology facilities [University of Manchester]. The histological slides were imaged and scored numerically using the Ashcroft scoring system by two experienced researchers blinded to the treatment groups. For rescue studies, 10 mg/kg SFN [LKT Laboratories] or 10 mg/kg vehicle [saline] was administered intraperitoneally at ZT6 prior to a bleomycin challenge at ZT12 and then every other day for 7 d before tissue processing.

Immunocytochemistry and immunohistochemistry

Immunostaining was performed as previously described [Pekovic et al. 2011]. Briefly, cells were fixed with 4% formaldehyde at 12-h intervals after stimulation with 50% horse serum. For immunocytochemistry, rabbit anti-NRF2 antibody [sc-722, Santa Cruz Biotechnology] was applied at 1:200 overnight at 4°C. Anti-rabbit IgG conjugated to Alexa Fluor 555 [Molecular Probes]. Nuclei were visualized by DAPI using VectaShield Hard-Set [Vector Laboratories]. Cells on cover slips were imaged using a Zeiss Axiovert 40 CFL inverted microscope [Carl Zeiss]. Intensity of NRF2 expression was measured using ImageJ software by measuring the integrated density of fluorescence in 50–100 cells after correcting for background fluorescence, and the results were confirmed in three independent experiments.

For immunohistochemistry, lung tissues were fixed with 4% formaldehyde, paraffin-embedded, and sectioned at 5 μm. Slides were dewaxed, rehydrated, and processed for antigen retrieval. Endogenous peroxidases were blocked in 3% H2O2 for 30 min. Blocking was carried out for 1 h in 2% normal goat serum, after which sections were incubated in anti-NRF2 antibody at 1:50 dilution overnight at 4°C in a humidified chamber. Slides were incubated in biotinylated secondary donkey anti-rabbit antibody [Vector Laboratories] diluted 1:800 in 1% BSA, ABC solution and the DAB [3,3’-diaminobenzidine] substrate kit [Vector Laboratories] were used to visualize positive NRF2 staining. To confirm the antibody specificity, immunohistochemistry was performed on control sections without the addition of either primary or secondary antibodies.

Statistical analyses

Data were evaluated using Student’s t-test, one-way ANOVA with Tukey test, or two-way ANOVA for multiple comparisons as indicated. Results are presented as mean ± SEM from at least
three independent experiments. Differences were considered significant at the values of $P < 0.05 (*)$, $P < 0.01 (**)$, and $P < 0.001 (***)$.

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