Tissue-specific and time-dependent regulation of the endothelin axis by the circadian clock protein Per1

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Abstract

Aims—The present study is designed to consider a role for the circadian clock protein Per1 in the regulation of the endothelin axis in mouse kidney, lung, liver and heart. Renal endothelin-1 (ET-1) is a regulator of the epithelial sodium channel (ENaC) and blood pressure (BP), via activation of both endothelin receptors, ET\textsubscript{A} and ET\textsubscript{B}. However, ET-1 mediates many complex events in other tissues.

Main methods—Tissues were collected in the middle of murine rest and active phases, at noon and midnight, respectively. ET-1, ET\textsubscript{A} and ET\textsubscript{B} mRNA expressions were measured in the lung, heart, liver, renal inner medulla and renal cortex of wild type and Per1 heterozygous mice using real-time quantitative RT-PCR.

Key findings—The effect of reduced Per1 expression on levels of mRNAs and the time-dependent regulation of expression of the endothelin axis genes appeared to be tissue-specific. In the renal inner medulla and the liver, ET\textsubscript{A} and ET\textsubscript{B} exhibited peaks of expression in opposite circadian phases. In contrast, expressions of ET-1, ET\textsubscript{A} and ET\textsubscript{B} in the lung did not appear to vary with time, but ET-1 expression was dramatically decreased in this tissue in Per1 heterozygous mice. Interestingly, ET-1 and ET\textsubscript{A}, but not ET\textsubscript{B}, were expressed in a time-dependent manner in the heart.

Significance—Per1 appears to regulate expression of the endothelin axis genes in a tissue-specific and time-dependent manner. These observations have important implications for our understanding of the best time of day to deliver endothelin receptor antagonists.

Keywords

Endothelin-A receptor; Endothelin-B receptor; ETA; ETB; Gene regulation
Introduction

The circadian clock regulates a variety of physiological processes such as metabolism, immune response, sleep–wake cycles, renal function, and blood pressure (BP) (reviewed in Richards and Gumz, 2012, 2013; Stow and Gumz, 2011). On the molecular level, the circadian clock consists of multiple proteins. Four are considered the core proteins that interact with one another to affect transcription of circadian target genes (Dibner et al., 2010). These proteins are Period (Per: homologs 1–3), Cryptochrome (Cry: homologs 1–2), BMAL1, and CLOCK. CLOCK and BMAL1 form a heterodimer, and then bind E-box DNA response elements to transcriptionally regulate CLOCK-controlled genes, including the genes Per and Cry. In the canonical model, Per and Cry presumably interact to repress the transcriptional activity of CLOCK and BMAL1 (Albrecht and Eichele, 2003).

Endothelin-1 (ET-1) is a peptide hormone expressed in multiple tissues and mediates its actions through two receptors: endothelin-A (ET_A) and endothelin-B (ET_B) receptors. ET-1 was first characterized as a potent vasoconstrictor; however, it is now known that ET-1 action is much more complex (reviewed in Kohan et al., 2011). ET-1 in the renal collecting duct is a potent inhibitor of epithelial Na channel (ENaC) activity through both ET_A and ET_B receptors (Bugaj et al., 2012; Ge et al., 2006, 2008; Lynch et al., 2013). This inhibition appears to occur via a nitric oxide-dependent mechanism (Bugaj et al., 2008; Stricklett et al., 2006) (reviewed in Kohan, 2013). The ET-1 gene (Edn1) is regulated by epigenetic factors (Welch et al., 2013) and transcription is controlled by mineralocorticoid action (Stow et al., 2009), calcium via the nuclear factor of activated T-cells (NFAT) (Strait et al., 2010), and a variety of other mechanisms (reviewed in Stow et al., 2011). Emerging evidence has demonstrated that Edn1 is also regulated post-transcriptionally (reviewed in Jacobs et al., 2013; Welch et al., 2013). Our laboratory has demonstrated that ET-1 peptide expression varies in a time-dependent manner in the renal cortex and medulla (Stow et al., 2012). The mechanism of this effect appears to be transcriptional. Indeed, we have previously shown that Per1 interacts with a non-canonical E-box from the Edn1 promoter (Stow et al., 2012). Per1 is a repressor of renal ET-1 mRNA and peptide levels, and Per1 knockout (KO) animals have elevated levels of ET-1 peptide in the kidney cortex and medulla (Richards et al., 2013; Stow et al., 2012).

Although ET-1 plays an integral role in a variety of physiological processes, circadian regulation of ET-1 and the receptors by the circadian clock and Per1 has not been studied outside of the kidney. Therefore, the goal of this study was to characterize the time-dependence of ET-1, ET_A and ET_B (the “endothelin axis”) mRNA expressions and to test the hypothesis that Per1 plays a role in the regulation of the endothelin axis mRNA in the liver, heart, kidney, and lung. It is well established that the circadian clock plays an integral role in the regulation of liver, heart, kidney and lung functions (reviewed in Richards and Gumz, 2013) but the role of Per1 in the regulation of the endothelin axis in these tissues has not been investigated. For the first time, we demonstrate that the endothelin axis is regulated by time and by Per1 in a manner that is unique to each of the tissues tested.
Materials and methods

Animals

All animal-use protocols were approved by the University of Florida and North Florida/South Georgia Veterans Administration Institutional Animal Care and Use Committee in accordance with the National Institutes of Health, Guide for the Care and Use of Laboratory Animals. Per1 KO and wild type (WT) mice (129/sv) were originally provided by Dr. David Weaver (University of Massachusetts (Bae et al., 2001)). WT and Per1 heterozygote (het) mice were bred in house by UF Animal Care Services Staff. Animals were maintained on a normal 12 h light: dark cycle. Mice were fed normal lab chow and given free access to water. At noon and midnight, mice were anesthetized and tissues were collected and snap frozen in liquid nitrogen. Kidneys were later dissected and cortex removed for protein or RNA isolation.

Isolation of IMCD

Isolations of inner medullary collecting ducts (IMCDs) were prepared as described below. WT and Per1 het mice were euthanized at midnight and inner medulla was dissected from both kidneys. The inner medulla was minced longitudinally and then digested at 37 °C in a buffer containing 250 mM sucrose, 10 mM Triethanolamine, 3 mg/mL of Collagenase type I, and 2 mg/mL of Hyaluronidase type IV for 30 min with gentle inversion. DNase I was then added at a concentration of 0.1 mg/mL and incubated for an additional 10 min. The mixture was then filtered over a 100 μm filter and the resulting supernatant spun at 600 x g for 3 min. The pellet was then suspended in a buffer containing 250 mM sucrose and 10 mM Triethanolamine and spun again. The pellet was then suspended in Hanks Buffered Salt Solution (HBSS—10 mM HEPES pH 7.4) and spun at 600 x g for 5 min.

Endothelin-1 ELISA

ET-1 peptide levels were determined as previously described (Stow et al., 2012). Cytosolic extracts were isolated using NE-PER kit (Pierce). Immunoreactive ET-1 peptide was detected by chemiluminescent ELISA (R&D Systems) and normalized to total protein content as determined by BCA assay (Pierce).

RNA isolation and qPCR

Total RNA was isolated using TRIzol (Invitrogen) according to the manufacturer’s instructions. RNA (10 μg) was treated with DNA-free DNase I (Ambion). DNaseI-treated RNA (2 μg) samples were used as template for reverse transcription with High Capacity cDNA Reverse Transcription Kit (Applied Biosystems). The resulting cDNAs (20 ng) were then used as template in quantitative real-time PCR (qPCR) reactions (Applied Biosystems) to evaluate changes in ET-1, ETA, and ETB mRNA levels. Cycle threshold (Ct) values were normalized against β-actin and relative quantification was performed using the ΔΔCt method (Livak and Schmittgen, 2001). Fold change values were calculated as the change in mRNA expression levels relative to the control. TaqMan primer/probe sets were purchased from Applied Biosystems.

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Statistical analysis

All data are represented as mean ± standard error of the mean (SEM). Statistics were performed with Graphpad Prism v6. All graphs and plots were made with Graphpad Prism v6. The effects of time and genotype were analyzed by two-way ANOVA with post-hoc Student–Newman–Keuls test. All *p* values less than 0.05 were considered significant.

Results

**Per1 regulates ET-1 expression but not ET<sub>A</sub> and ET<sub>B</sub> mRNA expression in the kidney**

We have previously shown that ET-1 peptide expression varies with time in murine renal cortex and inner medulla, with peak expression during the inactive period. In mice completely lacking the Per1 protein (Per1 knockout), renal ET-1 levels were increased (Stow et al., 2012). We have previously shown that Per1 heterozygous (het) mice have an approximate 50% reduction in Per1 protein expression in the kidney and liver (Richards et al., 2013) and that these mice exhibit a renal sodium wasting phenotype and reduced plasma aldosterone levels (Richards et al., 2013). To determine the effect of reduced Per1 expression on ET-1 expression, renal ET-1 peptide levels were measured in Per1 hetmice and compared to wild type (WT). ELISA was used to measure ET-1 peptide levels ex vivo in inner medullary collecting ducts (IMCD) isolated from WT and Per1 het mice at midnight. ET-1 peptide levels were significantly higher in IMCD from Per1 het mice than WT mice (Fig. 1).

To determine the effects of time and reduced Per1 protein levels on expression of the endothelin receptors, mRNA levels of ET<sub>A</sub> and ET<sub>B</sub> were assessed by qPCR in WT and Per1 het mice at noon and midnight, the middle of murine rest and active phases, respectively. In wild-type mice, ET<sub>A</sub> mRNA levels were significantly lower at midnight compared to noon in both the renal cortex and inner medulla (Fig. 2A), similar to the timed regulation of ET-1 peptide levels that we have previously observed (Stow et al., 2012). ET<sub>B</sub> mRNA levels also changed with time; however, the pattern was remarkably different between renal inner medulla and renal cortex (Fig. 2B). ET<sub>B</sub> mRNA levels were significantly higher at midnight in the renal inner medulla, but lower in the renal cortex at the same time. Per1 het mice had similar patterns of ET<sub>A</sub> and ET<sub>B</sub> mRNA expression with no apparent differences when compared to WT.

**Timed regulation and Per1-dependence of ET-1, ET<sub>A</sub>, and ET<sub>B</sub> mRNA expression in other tissues**

To examine timed regulation of the endothelin axis in non-renal tissues, mRNA levels of ET-1, ET<sub>A</sub>, and ET<sub>B</sub> were assessed in WT and Per1 het mice at noon and midnight in the heart, liver, and lung. In contrast to what was seen in the kidney, ET-1 mRNA levels in heart tissue from WT mice were significantly higher at midnight than at noon (Fig. 3A). Interestingly, the Per1 het mice exhibited elevated levels of ET-1 mRNA at both time points. ET<sub>A</sub> mRNA levels mirrored ET-1 mRNA in both genotypes (Fig. 3B). ET<sub>B</sub> mRNA levels remained constant at both time points for either genotype (Fig. 3C).
In the liver, ET-1 mRNA levels in WT mice were slightly higher at midnight than at noon (Fig. 4A). This variation in ET-1 mRNA levels with time was abolished in the Per1 het mice, but more dramatic effect was a substantial increase in ET-1 mRNA levels around the clock compared to WT. ET<sub>A</sub> mRNA levels were significantly lower at midnight in WT mice when compared to noon (Fig. 4B), and this effect was lost in Per1 het mice. Conversely, ET<sub>B</sub> mRNA levels were significantly higher at midnight when compared to noon in both genotypes (Fig. 4C).

In WT mice lungs, ET-1, ET<sub>A</sub> and ET<sub>B</sub> mRNA levels did not significantly change between noon and midnight (Fig. 5). Interestingly, Per1 het mice exhibited significantly lower ET-1 mRNA levels at both time points compared to WT mice (Fig. 5A), but no differences were observed in ET<sub>A</sub> and ET<sub>B</sub> mRNA levels (Fig. 5B and C). ET-1 peptide levels were measured in lung samples from WT and Per1 het mice euthanized at noon and midnight. ET-1 peptide levels were significantly lower in Per1 het mice at both time points (Fig. 5D) a result that corresponds to the changes observed at the mRNA level. Interestingly, WT ET-1 peptide levels in the lung increased significantly at midnight relative to noon.

**Tissue-specific differences in relative expression levels of ET<sub>A</sub> and ET<sub>B</sub> mRNA**

Both ET<sub>A</sub> and ET<sub>B</sub> are G protein-coupled receptors, but mediate distinct physiological responses. ET<sub>A</sub> is mostly associated with vasoconstriction, whereas ET<sub>B</sub> is normally associated with vasodilation (reviewed in Kohan et al., 2011). Therefore, the time-dependent relative levels of ET<sub>A</sub> and ET<sub>B</sub> would be informative for better understanding of ET-1 physiological action. We re-evaluated ET<sub>B</sub> mRNA expression relative to ET<sub>A</sub> in each tested tissue type to ascertain relative differences in receptor expression. In the renal inner medulla, mRNA levels of ET<sub>B</sub> were more than three-fold higher compared to ET<sub>A</sub> at noon for both genotypes and this difference jumped to six-fold at midnight (Fig. 6A). In the renal cortex at noon, ET<sub>B</sub> mRNA levels were only about two-fold higher than ET<sub>A</sub> (Fig. 6B). This difference was not apparent at midnight.

In contrast to the kidney, ET<sub>B</sub> mRNA expression in the heart is significantly lower than ET<sub>A</sub> at both noon and midnight (Fig. 7A). There did not appear to be an effect of reduced Per1 expression on the relative expression of the receptors in the heart. Similarly in the liver, significantly lower ET<sub>B</sub> mRNA levels were observed (Fig. 7B). However, this relative difference appeared to be Per1-dependent and was significantly decreased at midnight compared to noon. In the lung, there were no significant differences between ET<sub>A</sub> and ET<sub>B</sub> mRNA levels based on either time or genotype (Fig. 7C).

**Discussion**

The purpose of this study was to investigate the role of Per1 and time in the regulation of the endothelin axis in the kidney, heart, liver, and lung. Our investigation of mRNA expression in whole tissue samples clearly showed that tissue-specific and time-dependent regulation of ET-1, ET<sub>A</sub>, and ET<sub>B</sub> mRNAs occurs. We observed patterns of time-dependent and Per1-mediated regulation of the endothelin axis that were unique to each tissue type tested.
Recent work on the role of the endothelin axis in the kidney sheds light onto the novel function ET-1 plays in the regulation of sodium transport in the distal nephron. ET-1 is a potent inhibitor of ENaC activity via both the ET_A and ET_B receptors (Bugaj et al., 2012; Ge et al., 2008; Lynch et al., 2013), presumably through a nitric oxide-cyclic GMP-dependent mechanism (Bugaj et al., 2008; Stricklett et al., 2006). We have previously shown that ET-1 peptide levels vary in a time-dependent manner, peaking at noon, and that Per1 KO animals have increased ET-1 peptide in the renal cortex and inner medulla (Stow et al., 2012). In the present study, in both the renal cortex and inner medulla, ET_A mRNA was higher at noon than at midnight, coinciding with our previous report that ET-1 peptide levels were higher at noon than at midnight in mice (Stow et al., 2012). In contrast, ET_B mRNA levels exhibit opposite expression patterns in the inner medulla and cortex; ET_B mRNA levels are higher at midnight in the inner medulla, but lower in the cortex. However, ET_B and ET_A mRNA expression did not appear to be affected by reduced levels of Per1.

Although the receptors do not appear to be regulated in a Per1-dependent manner in this tissue, the timed regulation of ET_A and ET_B receptor expressions in the kidney should be investigated further to determine if there is a role for the ET receptors in the circadian control of sodium regulation.

ET_B mRNA levels are much higher than ET_A mRNA levels in both the renal inner medulla and cortex, and this observation is supported by previous reports using positron emission tomography which demonstrated that ET-1 binding to ET_B was significantly higher than ET_A binding in the rat kidney (Johnstrom et al., 2005). Also, a recent study demonstrated that renal collecting duct-specific ET_A KO mice have complete blunting of ET_A antagonist-mediated fluid retention (Stuart et al., 2013). This effect was not seen in cardiac-specific ET_A KO and was only partially evident in vascular smooth muscle-specific ET_A KO, confirming the tissue-specific expression/action of the receptor types.

In the heart, ET-1 and ET_A mRNA levels from WT mice were elevated at midnight as compared to noon. ET_A was found to be much higher than ET_B at all times. Activation of ET_A is primarily associated with increased contractility (MacCarthy et al., 2000), most likely through increases in intracellular calcium concentration (Talukder et al., 2001). It is possible that the increased ET_A levels at midnight act as a positive ionotrope for the increased activity during this time period.

ET-1 has also been shown to induce cardiac hypertrophy, presumably through an ERK1/2-dependent cascade (reviewed in Vignon-Zellweger et al., 2012). Recent findings have suggested that ET_B may play a role in this process as well. Cardiomyocyte-specific ET_A KO mice develop similar cardiac hypertrophy in response to angiotensin II infusion as WT controls (Kedzierski et al., 2003). Moreover, a recent study demonstrated that activation of the nuclear-membrane bound form of the ET_B receptor by intracellular ET-1 can result in inositol 1,4,5-triphosphate (IP_3) receptor-mediated calcium release, which is implicated in cardiachypertrophy (Merlen et al., 2013). Here ET_B mRNA was found to be constant and independent of circadian control. In contrast, both ET-1 and ET_A mRNAs were significantly increased in Per1 hetmice at both time points tested, suggesting loss of a negative regulatory mechanism. It should be noted that ET_A mRNA was significantly higher than ET_B mRNA expression, corroborating a previous study in rats (Fareh et al., 1996). These findings may
implies that an alteration in the ET<sub>A</sub>:ET<sub>B</sub> ratio, possibly through defects in negative regulatory mechanisms as in this study, or increased trafficking of the ET<sub>B</sub> receptor to the nuclear membrane could be involved in cardiac hypertrophy.

Much less is known about the physiological role of the endothelin axis in the liver. However, its role in liver pathogenesis is more apparent. ET-1 can promote activation of hepatic stellate cells (HSC), leading to increased cell proliferation, contraction, and survival (Pinzani et al., 1996; Rockey et al., 1998). HSCs, which express both ET<sub>A</sub> and ET<sub>B</sub> receptors, are pericytes found in the perisinusoidal space that once activated, lead to liver fibrosis and injury (reviewed in Yin et al., 2013; Housset et al., 1993; Pinzani et al., 1996). Angiotensin II has been shown to induce ET-1 production in HSCs (He et al., 1998). Another recent study demonstrated that interferon-gamma, a cytokine produced by T-cells that was previously shown to inhibit HSC proliferation and fibrogenesis, negatively regulates ET-1 expression (Li et al., 2012).

In normal liver function, ET-1 may act through an ET<sub>B</sub>-nitric oxide-dependent mechanism to increase expression of critical bile secretory genes and regulate choleresis, the secretion of bile into the gallbladder (Rodriguez et al., 2013). Here liver ET-1 mRNA increased somewhat at midnight and was substantially elevated in the livers of Per1 het mice at both time points. It will be interesting to see if Per1 knockout mice have an increased incidence of liver fibrosis or deregulation of bile secretion.

In the lung, it has been demonstrated that ET<sub>A</sub> receptor activation of pulmonary vascular smooth muscle cells can induce vasoconstriction through a mechanism involving IP<sub>3</sub> and calcium (Takwua et al., 1990), similar to what was observed in the heart. ET-1 was also linked to increasing proliferation of pulmonary vascular smooth muscle cells, through both the ET<sub>A</sub> and ET<sub>B</sub> receptors (Chua et al., 1992). However, deregulation of these processes contributes to pathophysiological states, including pulmonary arterial hypertension (PAH) (reviewed in Rubin, 2012). Patients with spontaneous PAH have increased levels of ET-1 in both serum and lung tissue compared to healthy controls (Giaid et al., 1993; Stewart et al., 1991). There are two commercially available endothelin receptor antagonists, ambrisentan and bosentan. Clinical trials have demonstrated the potential therapeutic benefits of these drugs in the treatment of PAH (reviewed in Seferian and Simonneau, 2013). Recently, a clinical trial demonstrated that a new dual ET<sub>A</sub> and ET<sub>B</sub> receptor antagonist, macitentan, led to significantly reduced morbidity and mortality among patients with pulmonary arterial hypertension (Pulido et al., 2013). Neither ET<sub>A</sub> nor ET<sub>B</sub> mRNA level varied with time of day in WT or Per1 het mice. However, the Per1 het mice did have significantly lower lung ET-1 mRNA expression at both noon and midnight, when compared to WT controls. Although time-dependent regulation was not apparent, reduced Per1 expression led to decreased ET-1 mRNA in the lung. It should be noted that this was a whole lung preparation. It would be interesting to examine the circadian pattern of ET-1 in just the pulmonary artery. Circadian regulation of the ET receptors in this tissue type could be important when timing drug administration for pulmonary arterial hypertension.

It is important to note that work over the last decade has begun to demonstrate potential “cross-talk” between the ET<sub>A</sub> and ET<sub>B</sub> receptors, which has arisen based on inhibitory...
studies that demonstrated inhibition of one of the receptors, which led to compensatory actions of the other (reviewed in Rapoport and Zuccarello, 2011). Multiple studies have shown the potential for ET$_A$ and ET$_B$ to hetero-dimerize, and that this interaction could be important for the observed “cross-talk” phenomena (reviewed in Watts, 2010). Relative receptor expression levels at a given time of day may affect the study of endothelin receptor cross-talk.

**Limitations**

The goal of this study was to determine the role of time and Per1 in the regulation of ET-1, ET$_A$ and ET$_B$ mRNA expression. With the exception of our demonstration that ET-1 peptide levels were significantly increased in IMCD cells but significantly decreased in lung tissue from Per1 het mice relative to WT control mice, our results are focused on changes in mRNA expression. Future studies are needed to determine if the effects of time and Per1 expression on the endothelin axis extend to the level of protein. Determination of relative protein expression at additional time points will be necessary in order to better understand the role of circadian rhythms in the regulation of ET-1, ET$_A$ and ET$_B$. Nevertheless, the results of the present study support the hypothesis that Per1 contributes to the regulation of the endothelin axis in a tissue-specific and time-dependent manner.

**Conclusions**

Results of the present study demonstrate that mRNA expression of the endothelin axis is regulated by Per1 in a tissue-specific and time-dependent manner. Presently, there is increasing interest in targeting the endothelin axis for the treatment of a variety of disorders, such as what is currently being done for the treatment of PAH. Multiple studies have demonstrated the benefit of chrono-pharmacotherapy, or timed administration of drugs to increase efficacy and decrease off-target effects. For example, it has been shown that chrono-pharmacotherapy in the treatment of hypertension leads to a decrease in adverse cardiovascular events (reviewed in Richards and Gumz, 2012). The results presented here suggest that optimization of the best time of day to deliver endothelin receptor antagonists should be considered.

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Fig. 1.
ET-1 peptide levels are elevated in IMCDs of mice with reduced levels of Per1. Wild type (WT) (light bars) and Per1 het (dark bars) mice were euthanized at midnight. IMCDs were isolated as described in the Materials and methods. ELISA was used to measure ET-1 peptide levels in WT and Per1 het mice. Data are presented relative to WT, ±SEM. ***P < 0.05 vs. genotype. N = 3.
Fig. 2.
Time-dependent regulation of ET<sub>A</sub> and ET<sub>B</sub> mRNA expressions in the renal inner medulla and cortex. Wild type (light bars) or Per1 het (dark bars) mice were euthanized at noon or midnight. Total RNA was isolated from dissected inner medulla and cortex and converted to cDNA. Real time quantitative RT-PCR (qPCR) was performed to evaluate mRNA expressions of ET<sub>A</sub> (Panel A) and ET<sub>B</sub> (Panel B). Data are presented relative to the WT at noon, ±SEM. †P < 0.05 vs. time, n= 3–4.
Fig. 3.
Regulation of ET-1, ETₐ and ET₉ mRNA expression in the heart by time and Per1. Wild type or Per1 hetmice were euthanized at noon or midnight. Total RNA was isolated from the heart and converted to cDNA. Real time quantitative RT-PCR (qPCR) was performed to evaluate in ET-1 (Panel A), ETₐ (Panel B), and ET₉ (Panel C). Data are presented relative to the WT at noon, ±SEM. *P < 0.05 vs. genotype; †P < 0.05 vs. time, n= 3–4.
Fig. 4.
Regulation of ET-1, ETA, and ETB mRNA expressions in the liver by time and Per1. Wild type or Per1 hetmice were euthanized at noon or midnight. Total RNA was isolated from the liver and converted to cDNA. Real time quantitative RT-PCR (QPCR) was performed to evaluate expression of ET-1 (Panel A), ETA (Panel B), and ETB (Panel C). Data are presented relative to the WT at noon, ±SEM. *P < 0.05 vs. genotype; †P < 0.05 vs. time; φ = significant interaction by 2-way ANOVA, n= 3–4.
Regulation of the endothelin axis in the lung by time and Per1. Wild type or Per1 het mice were euthanized at noon or midnight. Total RNA was isolated from the lung and converted to cDNA. Real time quantitative RT-PCR (qPCR) was performed to evaluate expressions of ET-1 (Panel A), ET\textsubscript{A} (Panel B), and ET\textsubscript{B} (Panel C). Data are presented relative to the WT at noon, ±SEM. *P < 0.05 vs. genotype; †P < 0.05 vs. time, n=3–4 (Panel D). ELISA was used to measure ET-1 peptide levels in the lung of WT and Per1 het mice at noon and midnight ±SEM. Data were analyzed using 2-way ANOVA, with significant genotype (*P < 0.05) and time (†P < 0.05) effects. There was no significant interaction between genotype and time. N = 3.

Fig. 5.
Fig. 6.
Relative differences between ET\textsubscript{A} and ET\textsubscript{B} mRNA expressions in renal inner medulla and cortex. Wild type or Per1 het mice were euthanized at noon or midnight. RNA was extracted and processed as described in Fig. 1. Real time quantitative RT-PCR (qPCR) was performed to evaluate relative differences in ET\textsubscript{A} and ET\textsubscript{B} expressions in renal inner medulla (Panel A) and cortex (Panel B). Data are presented relative to the WT ET\textsubscript{A} at noon, ±SEM. **p < 0.01, ***p < 0.001 vs. ET\textsubscript{A}; †P < 0.05 vs. time, n= 3–4.
Fig. 7.
Relative differences between ET_A and ET_B mRNA expressions in the heart, liver and lung. Wild type or Per1 hetmice were euthanized at noon or midnight. RNA was extracted and processed as described in Figs. 2–4. Real time quantitative RT-PCR (qPCR) was performed to evaluate relative differences in ET_A and ET_B expressions in the heart (Panel A), liver (Panel B), and lung (Panel C). Data are presented relative to the WT ET_A at noon, ±SEM. *P < 0.05, **p < 0.01 vs. ET_A; †P < 0.05 vs. time; P < 0.05 vs. genotype n = 3–4.