Diurnal profiling of neuroendocrine genes in murine heart, and shift in proopiomelanocortin gene expression with pressure-overload cardiac hypertrophy

Jennifer A Chalmers1*, Shuo-Yen J Lin1*, Tami A Martino1,3, Sara Arab3, Peter Liu1,4, Mansoor Husain1,3, Michael J Sole1,3 and Denise D Belsham1,2,3

Departments of 1Physiology and 2Medicine, and Obstetrics/Gynaecology, University of Toronto, Medical Sciences Building 3247A, 1 King’s College Circle, Toronto, Ontario, M5S 1A8 Canada
3Heart and Stroke/Richard Lewar Centre of Excellence for Cardiovascular Research and Toronto General Hospital Research Institute, Toronto, Canada
4Institute of Circulatory and Respiratory Health, Canadian Institutes of Health Research, Ottawa, Canada

(Correspondence should be addressed to D D Belsham; Email: d.belsham@utoronto.ca)

*J A Chalmers and S-Y J Lin contributed equally to this work

Abstract

Neuroendocrine peptides express biologic activity relevant to the cardiovascular system, including regulating heart rate and blood pressure, though little is known about the mechanisms involved. Here, we investigated neuroendocrine gene expression underlying diurnal physiology of the heart. We first used microarray and RT-PCR analysis and demonstrate the simultaneous expression of neuroendocrine genes in normal murine heart, including POMC, GnRH, neuropeptide Y, leptin receptor, GH-releasing hormone, cocaine- and amphetamine-regulated transcript, proglucagon, and galanin. We examined diurnal gene expression profiles, with cosinar bioinformatics to evaluate statistically significant rhythms. The POMC gene exhibits a day/night, circadian or diurnal, pattern of expression in heart, and we postulated that this may be important to cardiac growth and renewal. POMC diurnal gene rhythmicity is altered in pressure-overload cardiac hypertrophy, when compared with control heart, and levels increased at the dark-to-light transition times. These findings are also consistent with the proposal that neuropeptides mediate adverse remodeling processes, such as occur in pathologic hypertrophy. To investigate cellular responses, we screened three cell lines representing fibroblasts, cardiac myocytes, and vascular smooth muscle cells (NIH3T3, heart line 1, and mouse vascular smooth muscle cell line 1 (Movas-1) respectively). POMC mRNA expression is the most notable in Movas-1 cells and, furthermore, exhibits rhythmicity with culture synchronization. Taken together, these results highlight the diverse neuroendocrine mRNA expression profiles in cardiovasculature, and provide a novel model vascular culture system to research the role these neuropeptides play in organ health, integrity, and disease.

Journal of Molecular Endocrinology (2008) 41, 117–124

Introduction

The heart as an endocrine organ produces the hormone proopiomelanocortin (POMC; Forman & Bagasra 1992). POMC production is of particular interest, because it encodes many biologically active peptides relevant to cardiovascular physiology, such as adrenocorticotropic hormone, β-endorphin, and α- and γ-melanocyte-stimulating hormones (MSHs) (Diz & Jacobowitz 1983, Saito et al. 1983, Ekman et al. 1989, Millington et al. 1993, 1999, Ni et al. 2003). However, little is known about POMC gene expression in the heart and mechanisms of regulation in health and disease.

Previous studies demonstrate that >10% of cardiac genes exhibit diurnal expression patterns underlying homeostasis and pathophysiology (Young et al. 2001b, Storch et al. 2002, Martino et al. 2004, 2007a). Rhythmic gene expression in the heart is believed to underlie key physiologic functions, including heart rate, blood pressure, myocardial contractile performance, carbohydrate oxidation, oxygen consumption (Muller et al. 1985, Young et al. 2001a), and contribute to the timing of the onset of acute cardiac events (e.g., myocardial infarction, sudden cardiac death; Muller et al. 1985, Haus et al. 1989, Brzezinski 1997, Charloux et al. 1999). In this study, we focused on POMC gene expression in normal murine heart and pressure-overload cardiac hypertrophy. In addition, we extended investigations toward additional neuroendocrine genes expressed in the vascular-derived mouse vascular smooth muscle cell line 1 (Movas-1) cells. We find that POMC, in particular, exhibited rhythmic gene cycling similar to what is observed in the heart. Overall, these studies provide a new insight into the endocrinology relevant to cardiovascular function, and further provide a novel cell
model for examining the regulation of these peptides. Ultimately, these findings may translate to the clinical setting, providing evidence for the importance of the maintenance of normal circadian rhythms, critical for heart health (Martino et al. 2008).

Materials and methods

Animals

All animal protocols were reviewed and approved by the University of Toronto Institutional Animal Care and Use Committee. Six-week-old male C57Bl/6 mice were entrained to a normal 12 h light:12 h darkness cycle (LD 12:12) for 2 weeks time. They were then subjected to surgery or sham and maintained under the normal LD 12:12 cycle for a further 4 weeks. Animals were killed by decapitation and heart tissue was collected across the 24-h diurnal cycles, as described (Martino et al. 2007a). In a separate experiment to investigate gene expression in heart disease, pressure-overload cardiac hypertrophy was induced using a well-documented model of transthoracic aortic constriction (TAC) and compared with control (sham). Hearts from TAC and sham-operated mice were collected across the diurnal cycle, as described previously. A total of three samples were collected at each time point, at 4-h intervals over 24 h, from TAC and sham mice, and frozen at −80 °C until used.

Microarrays

Microarrays were used for global gene expression analysis, and we focused on the expression of neuroendocrine genes in the heart. We used the Affymetrix murine GeneChips (M0E430A). Briefly, RNA was purified from heart tissue by TRIzol reagent (Invitrogen). Integrity and highest quality were validated by ethidium bromide agarose gel electrophoresis, and BioAgilent spectrophotometry. First-strand cDNA synthesis, labeling, and hybridization to the microarray gene chips were performed using automated Affymetrix protocols. Scanning and per-chip normalizations were performed using MicroArray Suite version 5 (MASv5) and manufacturer specifications. GeneSpring (Agilent, Santa Clara, CA, USA) was used for visualizing gene expression patterns, and also for hierarchical cluster analyses, correlations, and ontology. Twenty-four hours of rhythmic gene profiles were determined by COSOPT analysis, a well-documented biocomputational program that interrogates daily gene cycling in microarray data (Panda et al. 2002, Martino et al. 2004). COSOPT provides an integer termed parameter multiple measures corrected-β (pMMC-β), which denotes the statistical significance of rhythms identified.

Movas-1

Movas-1 cells were grown in Dulbecco’s modified Eagle medium (DMEM; Sigma–Aldrich) supplemented with both 10% fetal bovine serum (FBS; Hyclone; distributed by Fisher Scientific, Ottawa, Canada), and 1% penicillin–streptomycin by volume (Gibco, distributed by Invitrogen) (Afroze et al. 2003). The cells were maintained in a tissue culture incubator at 37 °C, 5% CO₂. NIH3T3 cells were grown in DMEM with 10% FBS, and 1% pen–strept as described previously. The heart line 1 (HL-1) cell line was grown in Claycomb Media, 10% FBS (by volume), with 100 μg/ml pen–strept, 0·1 mM norepinephrine, and 2 mM L-glutamine, according to the protocol previously described (Claycomb et al. 1998).

Analysis of POMC gene expression in Movas-1 cells

Movas-1 cells were plated to confluence in 60 mm² tissue culture dishes and grown to confluency overnight (12–16 h; Falcon, distributed by VWR, Mississauga, Canada). For control or baseline values, the cells were scraped from the plates using a plastic scraping tool, every 4 h for 24 h. RNA was isolated using the guanidium thiocyanate–phenol–chloroform extraction method (Chomczynski & Sacchi 1987), and quantified via spectrophotometry to determine concentration and demonstrate high-quality purification. For experiments examining gene rhythmicity in vitro, confluent cells were first starved in serum-free media (supplemented with 5% pen–strept) for 12 h. The cells were then serum shocked with 20% FBS for 30 min. Serum shock is currently the standard method used to synchronize cells in culture (Hurst et al. 2002, Young 2006). Following serum shock, the cells were returned to normal culture conditions and samples were collected every 4 h for 36 h. Expression was determined by RT-PCR, as opposed to microarray.

Semi-quantitative reverse transcription (RT)-PCR

Gene expression patterns in the microarray data, and the Movas-1 cells, were both further confirmed by RT-PCR using the One-Step RT-PCR kit (Qiagen) according to the manufacturer instructions (see Table 1 for primer sequences). All PCR-amplified products were visualized on 2% agarose gels containing ethidium bromide, under u.v. light. Relative gene expression levels were quantified by densitometry on a Kodak Image Station 2000R. Expression values were normalized to histone. The PCR amplicons were verified by purification (Qiagen MinElute PCR Purification Kit) and sequencing (The Centre for Applied Genomics, the Hospital for Sick Children, MaRS Centre, Toronto, Ontario, Canada). Results were
compared for sequence homology by BLAST analysis on the NCBI website. Specific changes in gene expression patterns between test groups were evaluated statistically by ANOVA, followed by Bonferroni’s multiple comparison test (Prism 4.0). Statistical significance was determined at $P < 0.05$.

**Results**

**Neuroendocrine genes in the heart**

We first investigated expression of neuroendocrine genes in the normal murine heart. We detected expression of POMC, gonadotropin-releasing hormone (GnRH), neuropeptide Y (NPY), leptin receptor (ObRb), growth hormone-releasing hormone (GHRH), cocaine- and amphetamine-regulated transcript (CART), proglucagon (ProGlu), and galanin genes in the heart by microarray analysis, and subsequently verified these results using RT-PCR. Hypothalamic RNA was used as a control (Fig. 1).

**POMC gene expression in heart is circadian**

We examined the neuroendocrine genes for cosinear rhythms using COSOPT analysis, a well-developed biocomputational program that interrogates daily gene cycling in microarray data (Panda et al. 2002, Martino et al. 2004), in normal (sham-operated) mouse hearts. The POMC gene profile exhibited a robust rhythm over 24-h diurnal cycles, both by microarray analyses (Fig. 2A) and by semi-quantitative RT-PCR (Fig. 2B and C). The POMC gene expression peaked at night, ~4 h after lights off (ZT = 16), during the activity phase for mice (mice are active in the dark or nocturnal). The trough of POMC expression occurs during the day, ~4–7 h following lights on (ZT 4–7; subjective night or sleep).

**Aortic constriction alters POMC gene expression profile**

We next examined POMC gene expression in heart disease. We performed a parallel microarray study, but used mice subject to aortic constriction-induced cardiac hypertrophy versus sham-operated controls. As shown in Fig. 3, the sham mice exhibited the same expression profile as our control mice from the first experiment above, thus validating the methods used. Remarkably, POMC gene expression in TAC mice phase shifted relative to the sham control mice. That is, in contrast to normal heart, the peak for cardiac POMC expression in hypertrophied heart occurred ~8 h after lights off (ZT = 20). Additionally, significant differences were observed in the overall levels of POMC mRNA expression in cardiac hypertrophy, at ZT 3 and ZT 22 (*$P < 0.05$; Fig. 3).

<table>
<thead>
<tr>
<th>Gene</th>
<th>PCR template Heart Tissue</th>
<th>Cycle number</th>
<th>Movas-1</th>
<th>Annealing temp</th>
<th>Product size (bp)</th>
</tr>
</thead>
<tbody>
<tr>
<td>GnRH</td>
<td>S: actgtgtgttggagaagctgc A: ttcagatgtctctccagatc</td>
<td>40</td>
<td>40</td>
<td>62</td>
<td>162</td>
</tr>
<tr>
<td>GHRH</td>
<td>S: tgggtcctctcctctcagc A: atcactttgcccagctagag</td>
<td>40</td>
<td>40</td>
<td>60</td>
<td>128</td>
</tr>
<tr>
<td>ProGlu</td>
<td>S: ttcaccagcactacagcaga A: ggttgaatcagccagctgat</td>
<td>40</td>
<td>40</td>
<td>60</td>
<td>351</td>
</tr>
<tr>
<td>NPY</td>
<td>S: atgcacacagcgagaagcttc A: acatggaagggctctcaagc</td>
<td>40</td>
<td>40</td>
<td>60</td>
<td>282</td>
</tr>
<tr>
<td>POMC 1</td>
<td>S: tagatgtagtgactgctgtgc A: cagcagggggtgcgtctcagc</td>
<td>40</td>
<td>n/a</td>
<td>60</td>
<td>149</td>
</tr>
<tr>
<td>POMC 2</td>
<td>S: atgcccagatgtdtctagtcg A: tcacagtggctggcagaaaca</td>
<td>n/a</td>
<td>45</td>
<td>60</td>
<td>191</td>
</tr>
<tr>
<td>CART</td>
<td>S: atgctggcgctgctcctgcct A: cagctgccacagctcgcgtcctc</td>
<td>40</td>
<td>40</td>
<td>60</td>
<td>299</td>
</tr>
<tr>
<td>Galanin</td>
<td>S: cagcactggcatacagcaggagagctg</td>
<td>34</td>
<td>40</td>
<td>55</td>
<td>320</td>
</tr>
<tr>
<td>ObRb</td>
<td>S: atgcacacaggtactgctg A: ggcctcacttgcctgctgcctctg</td>
<td>28</td>
<td>40</td>
<td>60</td>
<td>356</td>
</tr>
<tr>
<td>Histone</td>
<td>S: gcagaggtgcctactgcctactg A: gggcctcacttgcctgctgcctg</td>
<td>21</td>
<td>25</td>
<td>60</td>
<td>217</td>
</tr>
</tbody>
</table>

S, sense; AS, antisense.
Expression of neuroendocrine genes in Movas-1 cells

We studied a pure heart-derived cell culture system, which can be used to examine diurnal neuroendocrine gene expression \textit{in vitro}. These cell lines do not have any sympathetic neuronal cell types that may be present when isolating the entire heart for analysis. We screened three cell lines representative of cardiac-derived cells: immortalized murine vascular smooth muscle cells (Movas-1), murine fibroblasts (NIH3T3), and cardiomyocytes (HL-1). We determined that only Movas-1 cells, similar to the cardiovasculature, express mRNA for \textit{POMC}, \textit{galanin}, \textit{ObRb}, tryptophan hydroxylase (\textit{TPH}), \textit{ProGlu}, and \textit{GnRH} (Table 2). It should be noted that the primer set used to amplify the \textit{POMC} mRNA in the Movas-1 cells detects the full-length \textit{POMC} mRNA and not the truncated form (Millington \textit{et al.} 1999). This full-length \textit{POMC} mRNA includes a region coding for a signal peptide that ensures translation of the transcript into protein. We then examined temporal gene expression of \textit{POMC} in the Movas-1 cells. The cells were synchronized by serum shock, consistent with methods for inducing circadian gene cycling \textit{in vitro} (Kaeffer & Pardini 2005). Movas-1 cells with serum shock showed a distinct pattern in \textit{POMC} expression, and this differed significantly from untreated controls at the 12- and 36-h time points ($P<0.001$ and $P<0.05$ respectively). When Movas-1 cells were treated with a 20% serum shock, \textit{POMC} mRNA exhibits a predictable pattern over time. This pattern in \textit{POMC} mRNA was nearly identical.

![Figure 1](image1.png)

\textbf{Figure 1} One-step RT-PCR screening results for neuropeptides in heart samples. Heart samples were positive for all neuropeptides tested. GnRH, gonadotropin-releasing hormone; GHRH, growth hormone-releasing hormone; NPY, neuropeptide Y; \textit{POMC}, proopiomelanocortin; CART, cocaine- and amphetamine-regulated transcript; ObRb, leptin receptor. Mouse hypothalamus was used as a positive control. M indicates molecular weight markers, NTC (no template control).

![Figure 2](image2.png)

\textbf{Figure 2} (A) Microarray data showing significant variation in relative \textit{POMC} mRNA expression over time in the normal murine heart. (B) One-step semi-quantitative RT-PCR conducted on heart tissue to confirm results from the microarray data, $n=3$ for each time point, for both banded and sham animals. (C) Representative gel of data shown in B, illustrating the change in relative \textit{POMC} mRNA expression compared with histone gene expression control over time. Animals were placed on a 12 h light:12 h darkness cycle (LD) schedule. Zeitgeber time corresponds to the time of day at which the tissue was harvested. Bar underneath graph indicates lights on in white and when lights were off in black.
to the rhythmic expression of the circadian clock gene brain and muscle aryl hydrocarbon receptor nuclear translocator (ARNT) like (BMAL1) that we have observed post-serum shock in the Movas-1 cells (JA Chalmers, TA Martino, N Tata, MR Ralph, MJ Sole & DD Belsham, unpublished observations), which suggests that they may be similarly regulated by the circadian clock. These results demonstrate synchronization of Movas-1 cells. Furthermore, cell culture synchrony by serum shock resulted in robust gene expression cycling in POMC mRNA, when compared with control.

Discussion

In this study, we describe a group of neuroendocrine genes expressed in murine heart, based on microarray and RT-PCR data. These included GHRH, CART, ProGlu, and galanin, which have never before been investigated in heart tissue, along with GnRH, NPY, POMC, and the long active form of ObRb mRNA. These findings add significantly to a growing list of neuroendocrine genes and products relevant to the cardiovascular system. We next examined the diurnal expression pattern of these genes in hearts from both control and TAC mice. We found that in the heart, the only gene with a distinct and significant diurnal rhythm was POMC (Supplemental Fig. 1, which can be viewed online at http://jme.endocrinology-journals.org/content/vol41/issue/). Therefore, we focused on this gene for the remainder of our study.

The POMC gene produces peptides previously localized to cardiocytes by immunohistochemical analyses (Diz & Jacobowitz 1983, Saito et al. 1983, Ekman et al. 1989, Millington et al. 1993, 1999, Ni et al. 2003). Here, we confirmed these findings and demonstrated that not only is it encoded by cardiac cells with a panel of other neuroendocrine genes, but also the POMC gene expression exhibited a rhythmic profile consistent with the 24-h period. We postulated that as such, POMC gene expression underlies physiology relevant to the cardiovascular system.

We next investigated POMC expression in vitro and provided evidence for a new model system for investigating relevant cell mechanisms. To do this, we screened several cell lines that represent different cell types found collectively in the heart. We examined the Movas-1 (vascular smooth muscle) cell line, the NIH3T3 (fibroblast) line, and the HL-1 (cardiomyocyte) line. Movas-1 was the only cell type to express POMC mRNA and, in addition, the cells were also positive for agouti-related peptide (AgRP), galanin, ObRb, TPH, proGlu, and GnRH. Furthermore, we demonstrated using serum-shock synchronization that POMC gene expression was rhythmic in Movas-1 cells. It is particularly interesting to observe rhythmicity in this key neuroendocrine gene outside of the brain. The profile of POMC expression in these cells closely mimics the pattern of BMAL1 (JA Chalmers, TA Martino, N Tata, MR Ralph, MJ Sole & DD Belsham, Unpublished observations). As BMAL1 gene expression in the brain and other tissues is regulated by several other circadian transcription factors, further work is required to first determine whether the clock

Figure 3 Microarray data displaying relative POMC mRNA expression over time from the heart tissue of mice that underwent TAC (Trans-aortic constriction) surgery versus sham animals over time. n=3 for each time point, for both banded and sham animals. POMC mRNA expression was significantly different in sham versus banded animals at points indicated (*P<0.05). The animals undergoing TAC showed a phase shift in peak POMC mRNA expression of 4 h.

Table 2: Results of one-step RT-PCR screening in mouse vascular smooth muscle cell line 1 (Movas-1) cell line for neuropeptides found to be present in the heart tissue

<table>
<thead>
<tr>
<th>Neuropeptides in Movas-1 cells</th>
</tr>
</thead>
<tbody>
<tr>
<td>Peptide</td>
</tr>
<tr>
<td>Cocaine- and amphetamine-regulated transcript (CART)</td>
</tr>
<tr>
<td>Galanin</td>
</tr>
<tr>
<td>Growth hormone-releasing hormone (GHRH)</td>
</tr>
<tr>
<td>Leptin receptor (ObRb)</td>
</tr>
<tr>
<td>Tryptophan hydroxylase (TPH)</td>
</tr>
<tr>
<td>Proglucagon (proGlu)</td>
</tr>
<tr>
<td>Proopiomelanocortin (POMC)</td>
</tr>
<tr>
<td>Gonadotropin-releasing hormone (GnRH)</td>
</tr>
<tr>
<td>Neuropeptide Y (NPY)</td>
</tr>
</tbody>
</table>

+, detectable; –, absent.

As culture synchronization is deemed important for coordinating cell cycle and/or circadian processes, this provides a potential new mechanism for understanding neuroendocrine gene expression and regulation in peripheral tissues in vivo.
directly controls POMC expression and, if so, which of these regulators is responsible (Reppert & Weaver 2001, 2002). The existence of E-box-binding sites in the promoter region of the POMC gene (Fig. 4) lends further support to the hypothesis of transcriptional regulatory control similar to that found in the brain and points to avenues of future direction (O’Donohue et al. 1979a, b, Monnet et al. 1981, Scimonelli et al. 1987).

Peripheral tissues, such as the heart, and also cell lines representative of non-neuronal tissues, express a remarkable collection of neuroendocrine genes. The regulation of these genes is likely important in organ structure and function, and further analysis of their role in cardiovascular endocrinology is warranted. For example, the peptide product γ-MSH, a natriuretic peptide derived from POMC, is actively investigated for its role in regulating a variety of cardiovascular functions, most notably affects on heart rate and blood pressure. This peptide is commonly thought to exert its cardiovascular action via the melanocortin 3 receptor, which binds with high selectivity and affinity in the anteroventral third ventricle region of the brain. We speculate that diurnal profiles of POMC mRNA synthesis, and peptide production, can facilitate the heart rate and blood pressure increase that occurs normally in the subjective day or active period, and the decrease at night or sleep period. Further support for this notion of diurnal availability of POMC for cardiovascular processes comes from findings that γ-MSH administered in the central nervous system (CNS) stimulates sympathetic outflow, increases mean arterial pressure, and produces tachycardia in most (Gruber & Callahan 1989, Wikberg et al. 2000, Schioth 2001), but not all (De Wildt et al. 1993, 1994) reports. Moreover, Ni et al. (2003) present evidence of a hypotensive role for γ-MSH in salt-sensitive hypertensive mice further supporting our hypothesis.

We observed a shift in the POMC diurnal gene expression profile in pressure-overload murine heart. The mechanism by which this occurs is not clear. One possibility is the underlying cell circadian mechanism. However, our previous studies demonstrated that the core clock genes (per2, bmal, rev-erb, etc.) and overall phase of rhythmicity in TAC heart do not change when compared with sham (Young et al. 2001b, Martino et al. 2007a). A more likely possibility is derived from the significant increase in blood pressure; this may contribute to the altered POMC expression profile in much the same manner as the matrix remodeling genes, which also change expression profiles in hypertrophic heart versus sham. Also, there is small vessel hyperplasia in heart along with perivascular fibrosis, a compensatory response to the high-pressure environment due to the aortic ligature. This observation of vasculopathy, in combination with the rhythmic expression of POMC gene expression in vascular myocytes (Movas-1) in vitro, provides a basis for the changes that occur. Lastly, there are very recent investigations by our group and others examining the diurnal proteome in peripheral organs (Reddy et al. 2006, Martino et al. 2007b). An interesting future experiment would be to determine which POMC-derived products are produced at differing times in the heart, and how these changes affect heart disease. Use of in vitro Langendorff could further provide evidence on the effects that occur in the day versus night, as well as responses to inhibitors given at specific times across the diurnal cycle.

In summary, we show expression of important neuroendocrine genes in the mammalian heart. Many of these gene products are involved in regulatory processes vital for cardiovascular physiology. In neural tissues, their timing of expression is presumed to underlie physiology, and thus we hypothesize a similar temporal role in peripheral tissues, such as the heart. We detect diurnal profiles of neuroendocrine genes expressed in normal heart. That diurnal profiles are also observed in heart disease, and specifically that POMC expression changes with the development of pathologic hypertrophy provides intriguing evidence for a potential role of circadian dysregulation in disease. We also investigated the non-neural cellular

![Figure 4](image-url)

**Figure 4** (A) Relative POMC mRNA expression in Movas-1 cells given no serum shock (dashed line) compared with cells given a 20% PFS serum shock (solid line) over time. Movas-1 cells displayed a distinct pattern over time after treatment with a serum shock, n= 3 with no serum shock, n= 4 with serum shock. Asterisks indicate differences between POMC mRNA expression with and without serum shock, *P* < 0.05, ***P*** < 0.001.

(B) Schematic of POMC promoter indicating the location of E-boxes where core clock transcription factors may potentially bind.
regulation of these genes, and demonstrated clear and reproducible expression in vascular myocytes in vitro. Culture synchronization by serum shock established a rhythmic gene phenotype in vitro and is consistent with neuroendocrine rhythms in the murine heart in vivo. These studies may shed new light and understanding on gene expression patterns underlying cardiovascular physiology and pathology and present novel avenues for future research in heart disease.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

Funding

This work was supported by the Heart and Stroke Foundation of Ontario (HSFO). D D B holds a Canada Research Chair in Neuroendocrinology and is a Canada Foundation for Innovation Researcher. J A C was funded by the National Sciences of Engineering Council.

Acknowledgements

T A M acknowledges the HSFO, and the HSRLCECR.

References


Hurst WJ, Mitchell JW & Gillette MU 2002 Synchronization and phase-resetting by glutamate of an immortalized SCN cell line. Biochemical and Biophysical Research Communications 298 133–143.


Journal of Molecular Endocrinology (2008) 41, 117–124

www.endocrinology-journals.org


Received in final form 28 May 2008

Accepted 11 June 2008

Made available online as an Accepted Preprint 12 June 2008