Thyroid hormone and vitamin D regulate VGF expression and promoter activity

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Abstract

The Siberian hamster (Phodopus sungorus) survives winter by decreasing food intake and catabolizing abdominal fat reserves, resulting in a sustained, profound loss of body weight. Hypothalamic tanycytes are pivotal for this process. In these cells, short-winter photoperiods upregulate deiodinase 3, an enzyme that regulates thyroid hormone availability, and downregulate genes encoding components of retinoic acid (RA) uptake and signaling. The aim of the current studies was to identify mechanisms by which seasonal changes in thyroid hormone and RA signaling from tanycytes might ultimately regulate appetite and energy expenditure. proVGF is one of the most abundant peptides in the mammalian brain, and studies have suggested a role for VGF-derived peptides in the photoperiodic regulation of body weight in the Siberian hamster. In silico studies identified possible thyroid and vitamin D response elements in the VGF promoter. Using the human neuroblastoma SH-SY5Y cell line, we demonstrate that RA increases endogenous VGF expression (P < 0.05) and VGF promoter activity (P < 0.0001). Similarly, treatment with 1,25-dihydroxyvitamin D3 increased endogenous VGF mRNA expression (P < 0.05) and VGF promoter activity (P < 0.0001), whereas triiodothyronine (T3) decreased both (P < 0.01 and P < 0.0001). Finally, intrahypothalamic administration of T3 blocked the short day-induced increase in VGF expression in the dorsomedial posterior arcuate nucleus of Siberian hamsters. Thus, we conclude that VGF expression is a likely target of photoperiod-induced changes in tanycyte-derived signals and is potentially a regulator of seasonal changes in appetite and energy expenditure.

Keywords

► VGF (non-acronymic)
► thyroid hormone
► SH-SY5Y cells
► Siberian hamster
► vitamin D

Introduction

The Siberian hamster (Phodopus sungorus) has been increasingly used to investigate hypothalamic mechanisms regulating energy homeostasis due to its natural seasonal cycle of appetite, energy expenditure, and body weight (Ebling 2014). These hamsters naturally become obese in the summer long-day photoperiod (LD), but then enter a catabolic state during winter short-day photoperiod (SD) where they reduce their food intake and catabolize intra-abdominal fat reserves, subsequently losing up to a third of their body weight (Bartness et al. 1989, Klingenspor et al. 1996, Mercer et al. 2001). The mechanism(s) by which these long-term changes in body weight are regulated are poorly understood, but they are clearly distinct from those governing short-term homeostatic regulation of energy balance.
appetite (Ebling 2015). The expression of a number of genes has been shown to be altered in the ventral ependymal wall lining the hypothalamic third ventricle of Siberian hamsters housed in different photoperiods (Barrett et al. 2005, 2006). Much recent interest has focused on tanycytes as the key component of the ependyma, as these cells are clearly important nutrient sensors in the hypothalamus, and also a stem cell niche (Bolborea & Dale 2013, Robins et al. 2013). For example, SD photoperiod upregulates expression of deiodinase 3 (DIO3), the enzyme responsible for degrading active 3,3',3'-triiodothyronine (T3) to inactive 3,3'-diiodothyronine (T2), as well as the conversion of thyroxine (T4) to the inactive 3,3',5-triiodothyronine, also called reverse T3 or rT3 (Barrett et al. 2005, 2006, 2007). A number of genes encoding components of retinoic acid (RA) uptake and signaling are also downregulated by SD in tanycytes (Rossi et al. 2004, Shearer et al. 2010). The major question now arises as to how seasonal changes in thyroid hormone and RA signaling from tanycytes to hypothalamic neurons ultimately regulate appetite and energy expenditure.

One of the most abundant peptidergic genes expressed in the mammalian brain, and particularly the hypothalamus, is VGF (non-acronymic) (Levi et al. 2004, Lewis et al. 2015b), a gene first identified on the basis of its rapid induction in vitro by nerve growth factor (NGF) in PC12 cells, a rat neuroblastoma cell line (Levi et al. 1985). It has subsequently been shown that RA, in addition to NGF, can act as a transcriptional inducer of the VGF gene in SK-N-NE (a human neuroblastoma cell line) and PC12 cells (Salton et al. 1991, Rossi et al. 1992, Cerchia et al. 2006). Hypothalamic VGF mRNA expression is altered by photoperiod in Siberian hamsters, with significantly lower expression in SD compared to LD in the arcuate nucleus (ARC), but intriguingly upregulation in a specific subdivision of the ARC, the dorsomedial posterior ARC (dmpARC), defined by expression of histamine 3 receptors (Barrett et al. 2005). After switching to LD, VGF expression in the dmpARC decreased rapidly, ahead of body weight changes (Barrett et al. 2005). We have previously shown that intracerebroventricular administration of the VGF-derived peptide, TLQP-21, decreased appetite and body weight in Siberian hamsters in LD (Jethwa et al. 2007), supporting the view that the products of this gene might impact upon seasonal behavior and physiology. In silico analysis of the mouse VGF promoter sequence revealed a potential thyroid response element (TRE), as well as a vitamin D response element (VDRE). We previously demonstrated that thyroid hormone (T3) availability in the hypothalamus was likely to be reduced in hamsters in SD, due to increase in DIO3 expression (Barrett et al. 2007, Murphy et al. 2012), while changes in vitamin D production have also previously been associated with photoperiod, particularly in the human kidney and skin (Webb et al. 1988, Holick 1995). Understanding the interactions of these regulatory factors is necessary to establish the mechanisms which promote the catabolic state observed in Siberian hamsters in SD. Thus, we investigated the effects of thyroid hormone (T3), RA and vitamin D (1,25-dihydroxyvitamin D3 [1,25D3]) on VGF mRNA expression and promoter activity in vitro. The experimental approach was to use the SH-SYSY neuroblastoma cell line, a common neuronal cell model due to its ability to differentiate into a more mature neuron-like phenotype and to be propagated unlike primary mammalian neurons (Dwane et al. 2013, Kovalevich & Langford 2013). These in vitro studies were complemented by an investigation of the effects of intra-hypothalamic implantation of T3 on the expression of VGF in the hypothalamus of Siberian hamsters, a procedure previously demonstrated to maintain an anabolic phenotype characteristic of LD exposure (Barrett et al. 2007, Murphy et al. 2012).

Methods

Materials

Unless stated otherwise, all chemicals for cell culture were purchased from Sigma–Aldrich, while those for RNA extraction, complimentary DNA (cDNA) synthesis and quantitative PCR (QPCR) were purchased from Roche Life Science. RA, T3, and 1,25D3 were obtained from Sigma–Aldrich and NGF was supplied by Millipore (Telecula, CA, USA) and was diluted as per manufacturer’s instructions. RA was reconstituted in 95% ethanol at 2.7 mg/ml; subsequent dilutions were made in DMEM with a final ethanol concentration of 0.1% (v/v). NGF (10 µg/ml) was reconstituted in sterile DMEM; subsequent dilutions were made in sterile DMEM. T3 was reconstituted in 1.0 ml 1.0 M NaOH (20 µg/ml) and 49 ml sterile DMEM; subsequent dilutions were made in DMEM. 1,25D3 was reconstituted in 95% ethanol (10 µM); subsequent dilutions were made in sterile DMEM.

Cell culture

The human neuroblastoma SH-SYSY cells (a kind gift from Dr Perry Barrett, but originally from European Collection of Cell Cultures (ECACC) Centre for Applied Microbiology and Research (CAMR), Porton Down Salisbury,
Wiltshire, UK) were grown in uncoated 25 cm² tissue culture flasks in DMEM/Ham’s F-12 containing 10% fetal bovine serum (FBS), 100 units/l penicillin and 100 mg/l streptomycin (DMEM/F12 complete) maintained at 37 °C in a 95% humidified incubator with 5% CO₂. Although SH-SY5Y cells tend not to adhere very well to uncoated plastic, they were routinely split 1:3 with 0.05% trypsin every 48 h.

**RNA extraction and cDNA synthesis**

Prior to harvesting the cells, the DMEM/F12 complete media was removed and cells were harvested in 200 µl of RNase-free PBS. Total RNA was extracted from the cells using the High Pure Isolation Kit (Roche) as described previously (Brown et al. 2012). First-strand cDNA was synthesized using the Transcriptor First Strand cDNA Synthesis Kit (Roche), according to the manufacturer’s protocol. The cDNA was stored at –20 °C.

**Quantitative RT-PCR**

The PCR was performed with SYBR green optimized for the LightCycler 480 (Roche Life Science). All reactions were performed in triplicate on 384 well plates as described previously (Brown et al. 2012). Transcript abundance was determined from a standard curve produced using a serial dilution of pooled cDNA made from all samples to check linearity and efficiency of the PCR and the values determined from a standard curve produced using a serial dilution of pooled cDNA made from all samples to check linearity and efficiency of the PCR and the values normalized to cyclophilin A, the most stable reference gene under the experimental conditions. The respective primer sets can be seen in Table 1.

**Synthesis of the VGF promoter and other constructs**

First, the ZsGreen gene present in the promoterless pZsGreen1-1 vector (Clontech Laboratory) was replaced with CBG992AmRFP from pCR2CBG992AmRFP (pRFP, Stritzker et al. 2014). The subcloning strategy utilized BamHI and NotI; then both fragments were purified using the QIAquick gel extraction kit (QIAGEN), as per manufacturer’s instructions. The pZsGreen1-1 backbone was treated with calf intestinal alkaline phosphatase (Promega) to prevent re-ligation of the plasmid without insert. This new reporter construct was designated pRFP-basic and was transformed into JM109 cells (Promega). Cultures were grown overnight, before plating and colony selection. All purified plasmids were subjected to restriction enzyme digestion and sequencing (performed by Source BioScience, Nottingham, UK) to confirm identity. Subsequently a cytomegalovirus (CMV) promoter, from pLenti6.4-CMV-C/EBPα (a kind gift from Prof Michael Lomax, University of Nottingham, UK), was cloned into the pRFP-basic vector. Using a similar strategy, pRFP-basic was digested with Xhol and SpeI, whilst pLenti6.4-CMV-C/EBPα was digested with SalI (which produces identical overhangs to Xhol) and SpeI (NEB, Hitchin, Hertfordshire, UK) to obtain the CMV promoter, which was subsequently ligated into the pRFP construct. This new construct was designated pCMV-RFP.

Approximately 1.1 kb of the mouse VGF promoter (−1151 bp to +51 bp, relative to the transcriptional start site; accession number: NM_001039385.1) was generated by PCR (see Table 1 for primers). The resultant amplicon was purified using the QIAquick gel extraction kit (QIAGEN), as per manufacturer’s instructions, and inserted into the pGEM-T-Easy vector for subsequent subcloning into pRFP-basic using SpeI and SacI. Alternatively, MboI and BglII were used to create the truncated 0.5 kb promoter construct lacking the potential TRE and VDRE. The orientation and authenticity of these constructs, designated pVGF1.1 and pVGF0.5, were verified by sequencing (performed by Source BioScience, UK).

**Study 1: the long-term maintenance of differentiation of SH-SYSY cells**

The SH-SYSY cells demonstrate reduced neurite length, a marker of differentiation, when cultured on uncoated surfaces (Dwane et al. 2013). Therefore to establish culture conditions for the long-term maintenance of differentiated neuronal cells, SH-SYSY cells were plated in

| Gene                  | Forward primer (5’–3’)               | Reverse primer (5’–3’)               |
|-----------------------|--------------------------------------|--------------------------------------|
| Cyclophilin A         | TCCGCCTTTCAGAATATTTC                 | ATTCGAGTTGTCACAGCTAGC                |
| Map2                  | CATGGTCTACAGGGCACCTATTCC             | GGTGGAAGAAGAGGCGAGATTACG            |
| Tau                   | GCCGGCGAGTGCTACATAGCTACA             | GGAAAGTCAGCTTGGGTTCCA               |
| Gap43                 | AGTAGGACAGGAGCAA                      | GTTGGGCAGGTAAGGT                |
| VGF                   | GACCCTCTCTCCACCTCTC                  | ACCGGCTCTTTATGTGCAGA              |
| VGF promoter          | AAGGGTGGGAGAGGAGTTG                  | GAGGGATGCGACCGGAG              |
six-well plates coated with or without 0.01% poly-1-lysine or 10 µg/ml collagen type IV. Cells were plated at $5 \times 10^4$ cells/cm$^2$ in DMEM/F12 complete, and 24 h later treated with 10 µM RA in DMEM/F12 complete for 120 h. Differentiation was subsequently maintained by treating the cells with 50 ng/ml NGF (Promega) in DMEM/F12 complete every 48 h. Images were captured 24, 48, 72, 96 and 120 h post-differentiation and neurite length determined using ImagePro Software (version 4.0; Image Pro, Rockville, MD, USA). A differentiated cell was defined as a cell with a neurite length greater than the length of the cell body. At the end of the study, cells were harvested for RNA and subsequent QPCR to determine the expression of the known markers of neurite differentiation, microtubule-associated protein 2 (Map2), Tau and growth-associated protein 43 (GAP43) (Table 1).

**Study 2: regulation of endogenous expression of VGF mRNA in vitro**

To investigate the regulation of endogenous VGF gene expression, undifferentiated SH-SYSY cells were plated onto uncoated six-well plates for 24 h. Subsequently, DMEM/F12 complete was removed and the cells treated with DMEM/F12 complete containing 50 ng/ml NGF, 10 µM RA, 10 nM T$_3$ or 10 nM 1,25D$_3$ for 24 h. For studies in differentiated SH-SYSY cells, plates were coated with 10 µg/ml collagen type IV and differentiated as per study 1. The adherent, differentiated cells were then treated with DMEM/F12 complete containing 50 ng/ml NGF, 10 µM RA, 10 nM T$_3$ or 10 nM 1,25D$_3$ for 24 h. Cells were harvested for RNA extraction and QPCR to determine endogenous expression of VGF.

**Study 3: regulation of VGF promoter activity in vitro**

To investigate the regulation of the VGF promoter in undifferentiated cells, SH-SYSY cells were plated onto uncoated six-well plates and grown to 70% confluence, prior to transfection in DMEM/F12 basic medium (DMEM/F12 containing 2.5% FBS, without antibiotics). The undifferentiated cells were transfected with the various plasmids using the FuGENE HD (reagent: DNA ratio of 3:1) as per manufacturer’s instructions. Briefly, undifferentiated cells were co-transfected with the VGF promoter construct (pVGF1.1 or pVGF0.5) and tenfold less pZsGreen1-N1 (Clontech Laboratories), a plasmid containing a variant of green fluorescent protein (GFP) under the control of a CMV promoter, with the latter used to correct for differences in transfection efficiency. pCMV-RFP (i.e. a strong promoter) was used as a positive control, while pRFP-basic (a promoterless plasmid) was used as a negative control, with both again being co-transfected with pZsGreen1-N1 to normalize for variability in transfection efficiencies. Seventy-two hours post-transfection, undifferentiated cells were switched back to DMEM/F12 complete (with antibiotics) containing 50 ng/ml NGF, 10 µM RA, 10 nM T$_3$, or 10 nM 1,25D$_3$ for 48 h. Promoter activities (fluorescence) were quantified at different time points using the Typhoon Trio+ (GE Healthcare, Little Chalfont, Buckinghamshire, UK).

In the experiments using differentiated cells, SH-SY5Y cells were grown to 70% confluence on uncoated six-well plates and transfected as before. The cells were then harvested using 0.05% trypsin, plated onto six-well plates coated with 10 µg/ml type IV collagen for 24 h, before differentiation was induced with 10 µM RA for 120 h (as before). Transfecting cells prior to differentiation have been shown to result in higher transfection efficiencies without altering the course of transgene expression (Lahousse et al. 2006, Chu et al. 2009). Transfected, differentiated cells were then treated with 50 ng/ml NGF, 10 µM RA, 10 nM T$_3$, or 10 nM 1,25D$_3$ for 48 h to investigate effects on the VGF promoter activities (up to 120 h post-transfection). Promoter activities (fluorescence) were quantified at different time points using the Typhoon Trio+ (GE Healthcare).

**Study 4: the effects of intra-hypothalamic T$_3$ administration on VGF mRNA expression in Siberian hamsters exposed to LD or SD**

Hypothalamic expression of VGF was studied in tissues collected in a study previously described by Barrett et al. (2007). The study was carried out in age-matched adult male Siberian hamsters obtained from a colony bred in house (Ebliing 1994), individually housed at constant temperature (21 ± 1°C) and 40–50% humidity. Animals had access ad libitum to food (Teklad 2019, Harland, UK) and water throughout the studies. All animal procedures were approved by the University of Nottingham Local Ethical Review Committee and were carried out in accordance with the UK Animals (Scientific Procedures) Act of 1986 (Project License PPL 40/2372).

Anesthetized Siberian hamsters (aged 3–4 months) maintained in LD had either T$_3$ (mixture of crystalline T$_3$ and medical grade silicone/Silastic-brand adhesive) or sham (medical grade silicone/Silastic-brand adhesive alone) microimplants placed bilaterally into the hypothalamus (6.5 mm below the surface of the dura at 0.5 mm...
to the left of the midline as defined by the center of the superior midsagittal sinus) as previously described (Barrett et al. 2007). At 12–16 days post-surgery, the Siberian hamsters were subdivided according to bodyweight to be either maintained in LD (n=6/group) or transferred into SD (n=7–8/group). Animals were euthanized with sodium pentobarbital (Euthatal: Rhone Merieux, Harlow) at 8 weeks post-surgery, and in situ hybridization studies for VGF were carried out as previously described in Barrett et al. (2005).

To determine expression slides were scored for the density of silver grains over individual cells in the dmpARC reflecting hybridization of the VGF probe by an observer who was blind to the treatment: 0 = no hybridization, 1 = a few cells expressing VGF mRNA, 2 = moderate VGF mRNA expression cells, 3 = abundant VGF mRNA. Scores were not possible for three animals as sections containing the dmpARC region were not available.

Statistical analysis
Data represent the means ± S.E.M. of six technical replicates (i.e. wells). Significant differences between groups for dependent variables were tested using either an unpaired, two-tailed Student’s t-test, one-way ANOVA in studies 1 and 2 or a two-way ANOVA (treatmentxtime) in study 3. In study 4, scores were analyzed by a Kruskal–Wallis test with post hoc Dunn’s tests for multiple comparisons. Changes in body weight (data represent the means ± S.E.M.) and in paired testis weights over the course of the T3 treatment were compared using one-way ANOVA with post hoc Dunnett’s tests for multiple comparisons. Statistical analyses were conducted using GraphPad PRISM (version 6.0, San Diego, CA, USA). Significance was accepted at P < 0.05.

Results
Study 1: the long-term maintenance of differentiation of SH-SY5Y cells
Treatment of the SH-SY5Y cells with 10 μM RA for 5 days significantly reduced proliferation (Fig. 1A and B, P < 0.01). The cells underwent a significant change in morphology, with the length of neurites significantly increasing in response to 10 μM RA (Fig. 1A and C, P < 0.0001). By coating the cell culture wells with an extracellular matrix protein such as poly-L-lysine or

![Figure 1](image_url)

Figure 1
Treatment of the SH-SY5Y cell line with 10 μM RA reduces cell proliferation and increases differentiation. (A) Images of SH-SY5Y cells at days 0 and 5 of differentiation in the absence or presence of 10 μM RA, showing differences in cell numbers and morphology (neurite lengths). (B) Treatment of the SH-SY5Y cell line with 10 μM RA significantly decreased cell number (P < 0.0001) and (C) significantly increased neurite length (a marker of differentiation) (P < 0.0001). (D) Neurite length was significantly greater 5 days post-treatment with differentiation media in wells coated with poly-lysine or type IV collagen (P < 0.0001) than plastic. (E) Treatment of the SH-SY5Y cell line with differentiation media significantly increased Map2, Tau and Gap43 (neuronal markers of differentiation). Gene expression was quantified by QPCR, normalized to cyclophilin A mRNA, and then compared to the normalized expression in undifferentiated cells. All values are means ± S.E.M. (n=6, *P < 0.05, **P < 0.01, ***P < 0.001, and **** P < 0.0001).
type IV collagen, neurite outgrowth was significantly enhanced (Fig. 1D, \( P < 0.0001 \)). Expression of Map2, Tau, and Gap43 mRNA were all significantly increased in comparison to undifferentiated controls (Fig. 1E, \( P < 0.05 \)). To ensure a homogenous population of differentiated SH-SY5Y cells, differentiated cells were then treated with 50 ng/ml NGF every 48 h. Under these conditions, cultures of differentiated cells could be maintained for up to 20 days without reversion or cell death.

**Study 2: regulation of endogenous expression of VGF mRNA in vitro**

Treatment of undifferentiated SH-SY5Y cells with 50 ng/ml NGF for 24 and 48 h significantly increased VGF mRNA five- and threefold (Fig. 2A, \( P < 0.01 \) and \( P < 0.05 \) vs vehicle treated controls), while treatment with 10 \( \mu \)M RA significantly increased VGF mRNA two- and fourfold at 24 and 48 h, respectively (Fig. 2A, \( P < 0.05 \) vs vehicle treated controls). Similar effects of NGF and RA were observed in differentiated SH-SY5Y cells, but the magnitude of the responses to NGF were bigger (Fig. 2B). Further studies showed that treatment of undifferentiated SH-SY5Y cells with 10 nM 1,25D3 for 24 h resulted in a threefold increase in VGF mRNA (\( P < 0.05 \)), whereas treatment with 10 nM T3 resulted in a fourfold decrease (\( P < 0.01 \)) in VGF mRNA (Fig. 2C), with similar effects again observed in differentiated SH-SY5Y cells, although the inhibitory effect of T3 tended to be greater (Fig. 2D).

**Study 3: regulation of VGF promoter activity in vitro**

Transfection of undifferentiated SH-SY5Y cells with pVGF0.5 or pVGF1.1 (containing 0.5 and 1.1 kb of the 5‘ flanking region of the VGF promoter respectively (Fig. 3A)) resulted in significant increases in fluorescence, indicating promoter activity, but there was no difference between them (\( P > 0.05 \), Fig. 3B). Treatment of transfected undifferentiated SH-SY5Y cells with either 10 \( \mu \)M RA or 50 ng/ml NGF resulted in significant increases in VGF promoter activities, with NGF inducing a much faster response than RA (time vs treatment interaction: \( F = 27.94, P < 0.0001 \), Fig. 3C). Similar time-dependent effects of both NGF and RA were observed in differentiated SH-SY5Y cells (time vs treatment interaction: \( F = 30.91, P < 0.0001 \), Fig. 3D). In both undifferentiated and differentiated SH-SY5Y cells, treatment with NGF resulted in a rapid induction of VGF promoter activity, whereas RA showed a much slower response with a delayed onset. Given the effect of T3 on endogenous VGF mRNA, transfected cells for promoter studies were pre-treated with 50 ng/ml NGF for 1 h (to briefly induce promoter activity), before removal and treatment with 10 nM T3. Treatment of undifferentiated or differentiated SH-SY5Y cells with 10 nM T3 significantly decreased promoter activity for the pVGF1.1 construct (time vs treatment interaction: \( F = 86.13, P < 0.0001 \), Fig. 3E and F), while treatment with 10 nM 1,25D3 resulted in a significant increase in pVGF1.1 promoter activity (time vs treatment interaction: \( F = 14.58, P < 0.0001 \), Fig. 3G and H). There were no effects (\( P > 0.05 \)) of either 10 nM T3 or 10 nM 1,25D3 on activity of the truncated plasmid (pVGF0.5), confirming that the response elements for the two nuclear receptors were only present in the longer promoter construct.

**Study 4: the effects of intra-hypothalamic T3 administration on VGF mRNA expression in Siberian hamsters exposed to LD or SD**

As previously reported (Barrett et al. 2007) exposure to SD for 8 weeks induced significant body weight loss (\( P < 0.01 \), \( P < 0.01 \)) for comparisons between control and treatment).
regulatory elements are present within the promoter 

as expected, a very low level of VGF expression was 

Promoter activities (fluorescence) were similar in cells transfected with 

UNet differentiation, cells were transfected as described in (B), but 72 h 

Differentiated transfected cells were then treated with 50 ng/ml NGF or 

increase in VGF abundance was not prevented by the 

Discussion

Tissue-specific expression of the VGF gene has been 

increase confirms the rise in VGF mRNA demonstrated in response to treatment with NGF in PC12 cells and to
RA in SK-N-BE cells (Levi et al. 1985, Cerchia et al. 2006). Whilst the increase in VGF mRNA and promoter activity in response to treatment with NGF was rapid, it was rather transient, decreasing after 24 h. In contrast, treatment with RA resulted in a much slower-, longer-term induction of VGF promoter activity, which continued to increase through to 48 h.

In silico studies identified possible TRE and VDRE sequences in the VGF promoter, and our studies have shown that both endogenous VGF mRNA expression and VGF promoter activity are suppressed by T₃ treatment, but increased with 1,25D₃. Correspondingly, an in vivo study revealed that intra-hypothalamic T₃ administration via slow-release microimplants reduced VGF mRNA expression in the dmpARC, arrow indicates induces expression in a SD sham hamster, RA-binding protein-2 (CRABP-2) and the nuclear receptors, RAR and RXR) is reduced in response to SD in the Siberian hamster (Ross et al. 2005, Barrett et al. 2006), so this may explain why VGF mRNA expression is reduced in the hypothalamus of Siberian hamsters exposed to SD (Barrett et al. 2005). Furthermore, in a photoperiod-responsive strain of rat, expression of RALDH-1, which converts retinol to RA, is also reduced in the hypothalamus in SD (Shearer et al. 2010). Transthyretin (TTR) is a transporter for vitamin A and its metabolite RA as well as T₄. TTR binds T₄ to establish a pool of T₄ in the plasma and cerebral spinal fluid (Prendergast et al. 2002) as well as transporting retinol by binding to the RBP (Hyung et al. 2010). However, studies utilizing TTR null mice have shown that while there are reductions in retinol and RBP in these mice, they display no symptoms of vitamin A deficiency, suggesting that TTR is not crucial for retinol delivery (Episkopou et al. 1993, van Bennekum et al. 2001). Thus, local availability of RA is determined by RALDH-1 and components of its signalling pathways (CRBP-1, CRABP-2 and RAR and RXR). All of which, as explained above, are reduced in response to SD in the Siberian hamster (Ross et al. 2005, Barrett et al. 2006, Shearer et al. 2010). Indeed, TTR expression in the hypothalamus of Siberian hamsters has been reported and the responsiveness of the gene was limited to the
photorefractory state (Prendergast et al. 2002). TTR mRNA is strongly expressed in the ependymal layer of the third ventricle and is decreased in SD (relative to LD) (Helfer et al. 2012), thus resulting in reductions in local T₃ availability. This in combination with the reduction in mRNA expression of DIO2 and increased expression and activity of DIO3 is observed in Siberian hamsters maintained in SD (Watanabe et al. 2004, Barrett et al. 2007, Herwig et al. 2009), and this results in a decrease in T₃. However, in the absence of RA, we suggest VGF expression would remain low in the hypothalamus of hamsters maintained in SD; thus we propose reduced RA in the hypothalamus of Siberian hamsters in response to SD results in reduced VGF expression (as demonstrated by Barrett et al. (2005)).

Previously, both vitamin A and RXR ligands have been shown to influence appetite (Anzano et al. 1979, Ogilvie et al. 2004). Therefore, the reduction in the availability of RA and components of its signaling pathway in the hypothalamus of Siberian hamsters in SD and subsequent reduction in VGF expression in the hypothalamus is a possible explanation for the effects on appetite. The complexity of the hormonal regulation of VGF expression that the current study has revealed may also explain why in SD there is a local upregulation of VGF expression in the dmpARC. In the hamster, this region has a much higher level of expression of thyroid hormone receptor b1 than the surrounding hypothalamus (Barrett et al. 2007), and our current study demonstrates that VGF mRNA expression in the dmpARC is specifically regulated by thyroid hormone. Thus, the SD-induced increase in DIO3 expression in tanyocytes would be expected to reduce local T₃ availability, resulting in a loss of repression of VGF gene expression in the dmpARC. Previously we demonstrated effects of increasing hypothalamic T₃ availability on the behavior and physiology of SD-exposed Siberian hamsters. Intra-hypothalamic T₃ implants placed in hamsters in SD produced a rapid increase in body weight, a reflection of increased food intake and a decrease in energy expenditure (Murphy et al. 2012). Here, we demonstrate that locally increasing hypothalamic T₃ blocks the SD-induced increase in VGF mRNA expression in Siberian hamsters. This is associated with a blockade of the SD-induced decrease in appetite and in weight loss, and also with the SD-induced inactivation of the reproductive axis. The correlation between VGF expression in the dmpARC and the physiological responses to SD was particularly highlighted in one individual hamster where the intrahypothalamic T₃ implants were ineffective in preventing any of the SD responses, probably because their placement was too rostral to influence hypothalamic function. The question now arises as to the specific role of T₃-regulated VGF expression in driving these seasonal responses. The function of the dmpARC itself is not clear; as one recent study found that SD-induced weight loss could occur in hamsters with lesions of this structure (Teubner et al. 2015). However, other lines of evidence suggest that increased VGF expression in the dmpARC could contribute to the SD-catabolic state, for example, at least one of the peptide products (TLQP-21) has been shown to reduce appetite when infused centrally into the hamster (Jethwa et al. 2007).

Repression of VGF promoter activity was nullified via the removal of the potential TRE from the promoter construct. TREs have previously been shown to be responsible for the dose-dependent T₃ repression of Mc4r promoter activity (Decherf et al. 2010). Furthermore, TREs have been shown to function in combination with RAR/RXR (De Luca 1991). Heterodimerization of the TR with RXR favors the dissociation of suppressors and the recruitment of activators of transcription (Cheung et al. 2009). Therefore, we hypothesize that the inability of TR to heterodimerize with RXR results in the repression of VGF in the hypothalamus of SD Siberian hamsters.

Additionally, we demonstrate that 1,25D₃, the active metabolite of vitamin D, significantly increases VGF endogenous mRNA expression and promoter activity in both undifferentiated and differentiated SH-SYSY cells. However, further studies are required to determine the effects of 1,25D₃ in vivo. The seasonal regulation of the hardierian gland of Siberian hamsters has been shown to be regulated by vitamin D (Perez-Delgado et al. 1993, Stumpf et al. 1993), while we have shown that plasma vitamin D₃ levels in adult hamsters are significantly higher in SD than in LD (SI Anderson, M Smith & FJP Ebling, unpublished observations). Interestingly, 1,25D₃ has been shown to have neuroprotective qualities in vitro (Brewer et al. 2001, Oermann et al. 2004, Wang et al. 2004), while treatment of the SH-SYSY cell line with 1,25D₃ inhibits proliferation (Celli et al. 1999), similar to the effects of RA. However, long-term incubation with 1,25D₃ only resulted in a slight trend towards differentiation (Celli et al. 1999). More recently, Agholme et al. (2010) demonstrated that pretreatment with RA followed by extracellular matrix gel adhesion, in combination with brain-derived neurotrophic factor (BDNF), neuregulin B1, NGF and 1,25D₃, resulted in differentiated SH-SYSY cells with unambiguous resemblance to adult neurons. The results presented here support the idea of differentiation-induced expression of the VGF gene and, therefore, a possible role in...
neurogenesis. This agrees with previous studies from Esposito et al. (2008), who suggests that receptor tyrosine kinase (RET) activation is a critical step in differentiation. Indeed, Korecka et al. (2013) have shown that RA induces RET expression in SH-SY5Y cells, while Cerchia et al. (2006) have suggested that inhibition of RET in SK-N-BE cells increases VGF expression. Moreover, a recent study in mice has demonstrated that the VGF-derived peptide, TLQP-62, directly increased the generation of neural progenitor cells in the hippocampus and potentiated BDNF-TrkB signalling (Thakker-Varia et al. 2014), thus the concept that enhanced VGF expression promotes neurogenesis and plasticity is rapidly gaining ground.

In conclusion, this study demonstrates that T<sub>3</sub> and 1,25D<sub>3</sub>, as well as NGF and RA, regulate endogenous VGF expression and promoter activity in vitro, while T<sub>3</sub> regulates VGF mRNA expression in vivo, providing a possible mechanism for the seasonal regulation of appetite in the Siberian hamster (Fig. 5), as well as suggesting a possible role for VGF in neurogenesis. Furthermore, it substantiates the central role of T<sub>3</sub> and 1,25D<sub>3</sub> in neuroendocrine and metabolic signaling.
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