EXPERIMENTAL STUDY

GH and IGF-I regulate the expression of endothelial nitric oxide synthase (eNOS) in cardiovascular tissues of hypophysectomized female rats

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Abstract

Objective: This study explored whether short-term replacement therapy with growth hormone (GH) affects blood pressure (BP), heart rate (HR) and endothelial nitric oxide synthase (eNOS) expression in cardiovascular tissues in hypophysectomized (Hx) female rats.

Design and Methods: BP, HR and the expression of eNOS in the aorta, caval vein and heart were studied in Hx female rats and in Hx female rats that underwent 7 days treatment with GH and thyroxine + glucocorticoids (T4 + GC). Insulin-like growth factor-I (IGF-I) was included in a second experimental protocol to explore the indirect effect of GH. The expression and localisation of eNOS was analysed by immunoblotting and immunohistochemistry.

Results: Decreased BP (Hx 98 ± 1; Intact 129 ± 3 mmHg; P < 0.05), HR (Hx 297 ± 14; Intact 399 ± 31 beats/min; P < 0.05) and unchanged eNOS expression was demonstrated in Hx compared with intact rats. None of the hormones affected BP, but both GH and IGF-I increased HR compared with Hx rats (GH 358 ± 10; IGF-I 337 ± 7, Hx 306 ± 11 beats/min; P < 0.05). Replacement of GH, GH + T4 + GC and IGF-I resulted in an increased aortic eNOS expression (GH 161 ± 24, GH + T4 + GC 177 ± 25, IGF-I 153 ± 21, Hx 109 ± 7%, P < 0.05), whereas in caval vein only GH + T4 + GC affected eNOS expression. None of the hormones changed the level of eNOS in the heart. eNOS was localised in the intima layer of the aorta, whereas in the caval vein eNOS was localised in all cell layers.

Conclusions: These findings support the suggested positive role of GH in the regulation of the cardiovascular homeostasis. The observed up-regulation of eNOS, and presumably an increased NO bioavailability, may result in improved endothelial and cardiovascular function.

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Introduction

Growth hormone (GH) deficiency or hyposecretion of GH is associated with an increased risk for cardiovascular mortality (1). These patients demonstrate normal blood pressure, hypotension or hypertension (for review see (2)). They also have endothelial dysfunction (3) and decreased nitric oxide (NO) formation (4). Several studies have shown the clinical benefits of GH therapy on cardiovascular function in these patients, such as unchanged or decreased systolic and diastolic blood pressure (BP), increased cardiac output (CO), increased heart rate (HR) (4, 5), increased NO formation (4) and decreased vascular resistance (5).

Hypophysectomized (Hx) rats demonstrate lowered BP, CO and HR (6, 7), as well as endothelial dysfunction (8), decreased heart weight and reduced wall to lumen ratio in the vasculature (9) in comparison to intact rats. Chronic treatment with GH results in improved vascular function (10). It has also been demonstrated that substitution with thyroxine (T4) and GH is required for the increase in BP and for the normal adaptation of the structure of the heart and the vessels in Hx rats (9).

It has previously been shown that formation of NO can be regulated via the GH–insulin-like growth factor-I (IGF-I) system (4). However, the cellular mechanism for this regulation is unknown. Nitric oxide (NO), formed by endothelial nitric oxide synthase (eNOS), has been shown to play an important role in the physiological regulation of local blood flow and BP (11, 12). Although considered to be constitutively expressed, different physiological factors are involved in the regulation of eNOS and NO formation (11).

The clinical consequence of hyposecretion of GH as well as replacement of GH for the cardiovascular function of the patient is unknown (13). The results of this study suggest improved cardiovascular function.
Materials and methods

Animals

Female Wistar rats were obtained from M&B (Ejby, Denmark). The animals underwent hypophysectomy at the age of 7 weeks (weight approximately 180 g) at M&B. All rats were acclimatized for 1 week before the onset of the experiment. All rats were weighed daily. None of the Hx rats increased their body weight (BW) prior to treatment (experiment (Exp) 1) from 162±1 to 163±5 g, Exp 2 from 170±1 to 169±1 g), demonstrating complete hypophysectomy (13), while the intact rats (Exp 1) demonstrated an increase in BW of ~13% (from 190±2 to 214±4 g). The rats had free access to standard pellet chow and tap water throughout the study. All experiments were approved by the Regional Animal Ethic Committee, Göteborg University.

Experiment 1

Hypophysectomised rats were randomised into four experimental groups: Hx (n = 10 rats), thyroxine + glucocorticoids (T4 + GC, n = 10), GH (n = 5) and GH + [T4 + GC] (n = 10). The hormones and vehicle were administered subcutaneously, twice daily for 7 days in a volume of 0.1 ml/100 g rats. Untreated intact rats (n = 10) were used as controls. The untreated Hx and intact rats received vehicle treatment.

Experiment 2

To explore whether the observed differences in central haemodynamics and aorta in Exp 1 were due to direct action of GH or mediated by IGF-I, IGF-I was included in the second experimental protocol. Three different experimental groups of Hx rats were used: Hx (n = 6), GH (n = 9) and IGF-I (n = 9). The supplementation of [T4 + GC] was not included in Exp 2, since no effects on either aortic eNOS expression in the cardiovascular tissues or on the BP and HR was detected after [T4 + GC] treatment in Exp 1. Growth hormone was administered subcutaneously twice daily, whereas IGF-I was delivered subcutaneously for seven days by a constant infusion using Alzet miniosmotic pumps (model 2001, Alza Pharmaceuticals, Palo Alto, CA, USA). These pumps were inserted into the neck during short-lasting barbiturate anaesthesia of the animals (methohexital sodium, Brietal; Eli-Lilly, Indianapolis, IN, USA) (75 mg/kg body weight, i.p.).

Measurement of plasma IGF-I

In intact rats (n = 7) from Exp 1 and untreated Hx rats (n = 6) and Hx rats treated for 7 days with either GH (n = 7) or IGF-I (n = 7) from Exp 2, the plasma IGF-I content was analysed by a commercial RIA kit (Mediagnost, Reutlingen, Germany). This RIA kit demonstrates high assay sensitivity as well as uses a highly specific polyclonal antibody that demonstrates about 49% in cross-reactivity to rat IGF-I (Mr Lindau, Mediagnost, personal communication). This suggests that the endogenous plasma IGF-I levels measured in all animals are underestimated by a factor of two.

Hormones (Exp 1 and 2)

The following hormones were used: thyroxine (L-thyroxine; Nycomed, Oslo, Norway) (T4, 10 μg/kg/day) and glucocorticoids (GC, cortisol phosphate; Solucortef, Upjohn, Puurs, Belgium) (400 μg/kg/day) (14). Bovine growth hormone (bGH, 1 mg/kg/day) (14) was generously supplied by American Cyanamid Co. (Princeton, NJ, USA). Insulin-like growth factor-I (IGF-I, 1.25 mg/kg/day) (15), was a generous gift from Genentech Inc. (San Francisco, CA, USA).

T4 and GC were diluted in saline, whereas bGH was diluted in 0.05 mol/l phosphate buffer (pH 8.6) containing 1.6% glycerol and 0.02% sodium azide.

Measurements of systolic blood pressure and heart rate (Exp 1)

Systolic blood pressure (SBP) and HR were measured by tail-cuff plethysmography (Narco BioSystems, Houston, TX, USA) on a randomised selection of rats from three of the groups in Exp 1: intact rats (n = 5), Hx rats treated with [T4 + GC] (n = 5) and Hx rats treated with GH + [T4 + GC] (n = 5). The measurements were done prior to hormone treatment (at days -7, -5 and -3), at the onset of treatment (day 0) and during treatment (at days 3, 5 and 7). Some of the SBP and HR data from intact rats and Hx treated rats with [T4 + GC] and GH + [T4 + GC] has been published before (16).

Measurement of conscious mean arterial blood pressure and heart rate (Exp 2)

Invasive conscious mean arterial blood pressure (MAP) and HR were measured for 30 min at day 7 in Hx (n = 5), GH-treated (n = 5) and IGF-I-treated (n = 5) rats. The procedure was performed by a catheterisation (PE-50) of the tail artery during short-lasting anaesthesia (methohexital sodium). The rats were allowed to
recover for at least 30 min after the operation before onset of registration.

**Dissection and homogenisation of tissues (Exp 1 and 2)**

The rats were killed by decapitation on day 7 of treatment. The inferior caval vein, aorta and heart were quickly excised. The heart was separated into left (including septum, LV) and right ventricles (RV). The tissues were trimmed free of fat and adherent tissues and weighed before being frozen in liquid nitrogen and stored at ~80°C until analysis.

Protein was extracted from the aorta, caval vein, LV, RV and liver as previously described (17). The total protein concentration was determined by a commercial protein assay (Bio-Rad, Hercules, CA, USA).

**Immunoblotting (Exp 1 and 2)**

The procedure for immunoblotting has previously been described in detail (18). Briefly, 25 μg (aorta, caval vein and liver) or 35 μg (LV, RV and liver) of total proteins were loaded in each lane on gels (10% NuPAGE Bis-Tris gels; Novex, San Diego, CA, USA). The gels were run for 90 min at constant voltage (150 V). Molecular weight standards (See Blue; Novex, San Diego, CA, USA) were used on each gel. The proteins were transferred to a polyvinylidifluoride (PVDF) membrane (Amersham, Buckinghamshire, UK). The membranes were then incubated with a mouse monoclonal antibody against eNOS (dilution 1:1000; Transduction Laboratories, Lexington, KY, USA). Immunoreactive protein was visualised by chemiluminescence using an alkaline phosphatase-conjugated secondary antibody (dilution 1:30,000; Sigma, St Louis, MO, USA) and CDP-Star (Tropix, Bedford, MA, USA) as a substrate. The membranes were exposed to ECL film (Amersham) at room temperature for 1–5 min and the films were subsequently developed. Semi-quantitative measurements of proteins from the immunoblots were made by densitometry (Fluor-S Multimager, Quantity One ver. 4.1.0, Bio-Rad). The optical density (OD) of each band was measured. The lane containing extract of liver was used as a reference on each gel. The lane containing protein was visualised by chemiluminescence using 

**Immunohistochemistry (Exp 1)**

Separate animals from the first experimental protocol (two in each treatment group: Intact, Hx, [T4 + GC], GH and GH + [T4 + GC]) were used for perfusion fixation for immunohistochemistry. Catheterisation of the common carotid artery with outflow from both jugular veins was performed during short-lasting barbiturate anaesthesia (methohexitol sodium). The rats were perfused at a pressure of 100 mmHg (Intact) or 80 mmHg (Hx) with phosphate-buffer (PBS) containing 0.58 M Na₂HPO₄, 0.17 M NaH₂PO₄ and 0.68 M NaCl for 15 min. The perfusion pressure was chosen in relation to the SBP in the respective groups (for details see Results and Fig. 1a). The solution was changed thereafter to 4% cold (+4°C) neutral buffered formaldehyde and the rats were perfused for an additional time of 30 min prior to the excision of the vessel. The aorta and caval vein were further fixated for 60 min in 4% formaldehyde and then transferred to 14% sucrose–PBS at 4°C for 24 h. The tissues were embedded in OCT-compound (Sakura Finetek Europe B.V., Zoeterwoude, the Netherlands) and snap frozen in 2-methylbutane chilled on dried ice, and stored at ~80°C until analysis (19).

The immunohistochemistry procedure has previously been described by Piontkewitz et al. (19). Briefly, the sections were blocked with 5% non-fat milk in PBS. This was followed by incubation with antibodies against eNOS (monoclonal antibody, dilution 1:50; Transduction Laboratories, Lexington, KY, USA), von Willenbrand factor (polyclonal antibody, dilution 1:800; Dako A/S, Glostrup, Denmark), or smooth muscle actin (monoclonal antibody, dilution 1:1600; Sigma). After several rinsing steps the sections were incubated with either biotinylated anti-mouse or biotinylated anti-rabbit (dilution 1:200 for both; Vector Laboratories, Burlingame, CA, USA). Rat serum (dilution 1:50) was added in conjunction with the anti-mouse in order to reduce unspecific binding. Thereafter, the sections were developed in streptavidin fluorescein (dilution 1:200; Amersham, Buckinghamshire, UK) and mounted with Moviol/Dabco mounting medium. The following controls for unspecific binding were used: replacement of primary antibody with (1) non-specific anti- IgG1 antibody (dilution 1:800, Dako A/S, Glostrup, Denmark); or smooth muscle actin (monoclonal antibody, dilution 1:1600; Sigma). After several rinsing steps the sections were incubated with either biotinylated anti-mouse or biotinylated anti-rabbit (dilution 1:200 for both; Vector Laboratories, Burlingame, CA, USA). Rat serum (dilution 1:50) was added in conjunction with the anti-mouse in order to reduce unspecific binding. Thereafter, the sections were developed in streptavidin fluorescein (dilution 1:200; Amersham, Buckinghamshire, UK) and mounted with Moviol/Dabco mounting medium. The following controls for unspecific binding were used: replacement of primary antibody with (1) non-specific anti- IgG1 antibody (dilution 1:800, Dako A/S, Glostrup, Denmark); or (2) 5% non-fat milk. These controls demonstrated negligible signals (data not shown).

**Statistical analysis**

Values are given as means±s.e.m. Left ventricular and RV weights (Exp 1), MAP and HR (Exp 2), changes in BW (Exp 1 and 2) and levels of plasma IGF-I were analysed by ANOVA followed by Fisher’s LSD as a post hoc test. ANOVA adapted for repeated measures was used to evaluate SBP and HR in Exp 1. The Mann–Whitney rank sum test was used for the analysis of immunoblotting data obtained by densitometric scanning from both experimental protocols. A P value of less than 0.05 was considered significant.

**Results**

**Experiment 1**

**Body weight** The BWs of Hx rats and Hx rats receiving [T4 + GC] were unchanged throughout the experiment (Hx from 164±1 to 161±1 g; [T4 + GC] from 160±3
to 157±4 g). The intact rats showed a slight, but non-significant, increase in BW at day 7, compared with day 0 (from 214±4 to 218±3 g). Rats supplemented with GH or GH+[T4+GC] demonstrated a significant increase in BW (GH from 161±2 to 183±4 g, P < 0.05; GH+[T4+GC] from 161±2 to 187±2 g, P < 0.05).

**Systolic blood pressure and heart rate** Hypophysectomy per se, resulted in a decrease in both SBP and in HR compared with intact animals (P < 0.05; Fig. 1A and 1B). During replacement (days 0–7), the SBP remained unchanged in all treatment groups (Fig. 1A). Heart rate remained unaltered after replacement with [T4+GC] in Hx rats (Fig. 1B). However, the animals receiving GH+[T4+GC] demonstrated an increase in HR from day 3 (P < 0.05 compared with day 0 and between [T4+GC] and GH+[T4+GC] during the treatment period) and reached the same HR level as intact animals (Fig. 1B). The HR remained elevated until the experiment was completed (Fig. 1B).

**Left and right ventricular weights** Hypophysectomy caused a significant decrease in LV (P < 0.05), but not in RV weight, compared with intact rats (Fig. 2A and B). After treatment, LV weight increased in [T4+GC], GH, and GH+[T4+GC] groups (P < 0.05 for all vs. Hx, Fig. 2A). There was a small, but significant difference between [T4+GC] and GH+[T4+GC] in LV weights (P < 0.05, Fig. 2A). None of the treatments had any effect on the RV weight (Fig. 2B).

**Expression and cellular localisation of eNOS in aorta, caval vein, RV and LV**

**Intact and Hx rats** There were no differences in the eNOS protein expression in the aorta, the caval vein (Figs 3, 4 and 5a–d), the RV (data not shown) or the LV (data not shown) between Hx rats and intact rats.

The expression of eNOS was higher (1.6-fold) in the caval vein compared with the aorta in Hx animals (P, 0.05; Figs 3 and 4). There was no difference in eNOS expression between LV and RV in Hx rats (data not shown).

**Figure 1** Systolic blood pressure (SBP, A) and heart rate (HR, B) prior to and during treatment. Measurements at -7, -5 and -3 days are pretreatment values. Treatment with different hormones (thyroxine + glucocorticoids ([T4+GC]) and growth hormone (GH)) were started on day 0. SBP and HR were measured at days 3, 5 and 7. The different groups were hypophysectomized (Hx) treated rats with [T4+GC] (n = 5) and GH+[T4+GC] (n = 5). Intact rats (n = 5) served as controls. Data is expressed as means±S.E.M. ANOVA adapted for repeated measures was used for statistical analysis. *P < 0.05 between intact control and Hx rats. †Significant change in GH+[T4+GC]-treated rats compared with day 0 (P < 0.05). ‡P < 0.05 between GH+[T4+GC] and [T4+GC] during the treatment periods (day 3, 5 and 7, P < 0.05). Some of the SBP and HR data from intact rats and Hx rats treated with [T4+GC] and GH+[T4+GC] has been published before (16).
The aorta showed a specific staining for eNOS in the endothelial cells (Fig. 5a and c). Unspecific staining of the elastine components of the media layer was observed in the aorta (Fig. 5a, c, e, g and i). This was also noticed with non-specific IgG1 (data not shown). In the caval vein, the staining included cells localised to the intima, media and adventitia layers (Fig. 5b and d).

\[ T_4 + GC \] - GH- and GH + [T₄ + GC]-treated Hx rats The expression of eNOS in aorta, caval vein, LV or RV was not affected by \[ T_4 + GC \] (Figs. 3 and 4). There was an increase of eNOS expression in the aorta after GH supplementation compared with Hx and Hx rats treated with \[ T_4 + GC \] (\( P < 0.05 \) for both, Fig. 3). This increase was also observed in the aorta when GH was combined with \[ T_4 + GC \] (\( P < 0.05 \), Fig. 3). The caval vein of GH + [T₄ + GC]-treated animals demonstrated a 1.4-fold higher eNOS expression compared with the aorta in the same animals (\( P < 0.05 \), Figs 3 and 4), which was similar a difference as seen in Hx rats. An increased eNOS expression was shown in the caval vein of GH + [T₄ + GC]-treated animals compared with \[ T_4 + GC \] rats (\( P < 0.05 \), Fig. 4). The hormonal

Figure 2 Left (A) and right (B) ventricular weights of intact (controls, \( n = 10 \)), hypophysectomized (Hx, \( n = 10 \)) and Hx rats treated thyroxine and glucocorticoids ([T₄ + GC]) (\( n = 10 \)), growth hormone (GH, \( n = 5 \)) or GH + [T₄ + GC] (\( n = 10 \)) for 7 days. Data is expressed as means ± S.E.M. Changes in left and right ventricular weight were analysed by ANOVA followed by Fisher’s LSD as a post hoc test. \( P < 0.05 \) between controls and Hx, \( P < 0.05 \) between Hx and the different treatments and \( P < 0.05 \) between [T₄ + GC] and GH + [T₄ + GC].

Figure 3 The protein expression of endothelial nitric oxide synthase (eNOS) in the aorta of intact rats (controls, \( n = 10 \)), hypophysectomized (Hx, \( n = 10 \)) and Hx rats after 7 days of treatment with thyroxine and glucocorticoids ([T₄ + GC]) (\( n = 10 \)), growth hormone (GH, \( n = 5 \), or GH + [T₄ + GC] (\( n = 10 \). The values are obtained by densitometric scanning of immunoblots. Data is expressed as means ± S.E.M. and as a percentage of expression in liver (100%). The same liver sample was used as the standard for each gel. The Mann–Whitney rank sum test was used for statistical analysis. \( P < 0.05 \) vs. intact, \( P < 0.05 \) vs Hx and \( P < 0.05 \) vs [T₄ + GC].
Immunohistochemistry showed specific eNOS staining in the endothelial cells in the intima layer in the aorta (Fig. 5e, g, and i). In the caval vein, the staining for eNOS was observed in endothelial cells of the intima-, media- and adventitia cell layers (Fig. 5f, h and j). No alteration in the cellular localisation in aorta or caval vein was observed after hormonal treatments compared with controls (intact, Hx) (Fig. 5).

The aorta demonstrated unspecific staining for eNOS of the elastine components of the media layer (Fig. 5a, c, e, g and i), which also was observed with non-specific IgG1 (data not shown).

Expression and cellular localisation of von Willenbrand factor and smooth muscle actin

In the aorta, only the endothelial cells in all groups demonstrated staining for von Willenbrand factor (Fig. 6a). In contrast, staining for this factor was present in all layers (intima, media and adventitia) of the caval vein (Fig. 6b). No difference in staining for von Willenbrand factor was observed between the different experimental groups.

The localisation of smooth muscle cells was demonstrated by staining of α-smooth muscle actin in both the aorta and the caval vein. All treatment groups showed the same specific staining of α-smooth muscle actin in the media of aorta and caval vein (Fig. 6c and d).
**Experiment 2**

**Body weight** In the second experimental protocol, the BW of Hx rats was unchanged throughout the study (from 173±2 to 173±1 g). GH treatment in Hx rats demonstrated increased BW from 168±2 to 190±3 g (P < 0.05) and after IGF-I supplementation the BW increased from 161±2 to 177±2 g (P < 0.05).

**Table 1** Plasma insulin-like growth factor-I (IGF-I) levels in intact rats, untreated hypophysectomized (Hx) rats and Hx rats treated with either growth hormone (GH) or IGF-I for 7 days. Values are expressed as means ± S.E.M.

<table>
<thead>
<tr>
<th>Group</th>
<th>N</th>
<th>Plasma IGF-I (ng/ml)</th>
</tr>
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<tbody>
<tr>
<td>Intact</td>
<td>7</td>
<td>491±55</td>
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| Hx    | 6  | 43±6                *
| GH    | 7  | 273±18*             |
| IGF-I | 7  | 319±26*             |

ANOVA followed by Fisher’s LSD as a post hoc test was used for counting statistically significant changes, *P < 0.05 vs intact, **P < 0.05 vs Hx.

**Plasma IGF-I** The Hx untreated rats showed a statistically significant decrease in plasma IGF-I compared with intact rats (Table 1, P < 0.05). There was a substantial increase of plasma IGF-I in GH- and IGF-I-treated Hx rats compared with untreated Hx rats (Table 1, P < 0.05), whereas both GH- and IGF-I-treated Hx rats demonstrated lower plasma IGF-I levels compared with intact rats (Table 1, P < 0.05). There was no significant difference in plasma IGF-I levels between GH- and IGF-I-treated Hx rats (Table 1).

The same pattern of IGF-I levels were found in intact, Hx and GH groups when compensating for the fact that the IGF-I antibody showed different cross-reactivity to human and rat IGF-I (P < 0.05). However, the levels of IGF-I were significantly lower in the IGF-I group compared with the GH group (P < 0.05). The recalculated levels were: Intact 981±110, Hx 85±12, GH 547±35 and IGF-I 362±26 ng/ml.

**Conscious mean arterial blood pressure and heart rate** No change in conscious mean arterial blood pressure (MAP) measurement was observed between the groups (Table 2). Furthermore, the GH-treated Hx rats showed a 17% increase in HR compared with untreated Hx rats measured by conscious HR measurement (P < 0.05, Table 2). An increase in HR was also demonstrated in IGF-I-treated Hx rats compared with untreated Hx rats (P < 0.05, Table 2).

**Table 2** Invasive conscious mean arterial blood pressure (MAP) and heart rate (HR) measurements in hypophysectomized (Hx) rats and Hx rats treated for 7 days with either growth hormone (GH) or insulin-like growth factor-I (IGF-I). Values are expressed as means ± S.E.M.

<table>
<thead>
<tr>
<th>Group</th>
<th>n</th>
<th>MAP (mmHg)</th>
<th>HR (beats/min)</th>
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<tr>
<td>Hx</td>
<td>5</td>
<td>93±3</td>
<td>306±11</td>
</tr>
<tr>
<td>GH</td>
<td>5</td>
<td>94±2</td>
<td>358±10*</td>
</tr>
<tr>
<td>IGF-I</td>
<td>5</td>
<td>98±2</td>
<td>337±7*</td>
</tr>
</tbody>
</table>

ANOVA followed by Fisher's LSD as a post hoc test was used for counting statistically significant changes between Hx and Hx-treated (GH or IGF-I) groups, *P < 0.05.
Data are expressed as means ± S.E.M. and as a percentage of normal LV weight in these rats, but it seems that longer replacement therapy is needed to create a normalisation of the vascular structure.

In the veins another indirect effect of GH may mediate the increased eNOS expression, but had no effect on the eNOS expression in the other studied tissues. In addition, the distribution of eNOS differed between the two vessel types. In the aorta, the expression of eNOS was localised to the endothelial cells of the intima, whereas the caval vein expressed eNOS in all cell layers (intima, media and adventitia). Our results suggest that the effect on eNOS expression in the arteries (aorta) could partly be mediated by IGF-I and/or by a direct effect of GH on the vascular cells. In the veins another indirect effect of GH may mediate this increased eNOS expression.

**Cardiovascular function**

The present study demonstrated decreased BP and HR after hypophysectomy. These results are in accordance with those of László et al. (6), Shen et al. (7) and Folkow et al. (9). A reduced structural adaptation of the left ventricle (present study and (9)) as well as lower CO (6, 7) together with loss of vascular structure, e.g. reduced synthesis of collagen (20) and reduced wall/lumen ratio in the resistance vessels (9), may explain the reduction of BP in Hx rats. After 7 days of supplementation with [T₄ + GC], GH or IGF-I the LV weight increased up to the levels of intact rats, although the BP was still reduced compared with intact rats in all hormonal treatment groups. Folkow et al. (9) demonstrated that 6 weeks of GH + T₄ supplementation to Hx rats restored the blood pressure, LV weight and the vascular structure to those of intact rats. Our study may suggest that 1 week of hormonal replacement seems to be enough to create normal LV weight in these rats, but it seems that longer replacement therapy is needed to create a normalisation of the vascular structure.

Both in the present study and in studies involving humans (4), an increase in HR has been demonstrated after supplementation with either GH or IGF-I. This finding (increased HR and unaltered BP) can be explained by: (i) a stimulatory effect of GH and IGF-I on the heart followed by dilatory response in the resistance vessels, or (ii) a primary effect (dilatation of vessels) of GH and IGF-I with a reflexogenic increase in HR to maintain BP. Our results support the second explanation since the major effect of GH and IGF-I on eNOS protein expression was observed in the aorta, leaving the level of eNOS protein in the heart unaltered. An increase in eNOS expression may result in an increased NO bioavailability, causing a decreased total peripheral resistance (TPR). A decreased TPR after GH supplementation to a GH-deficient patient has previously been reported (5). Thus, this may suggest that an increased eNOS expression in the arteries together with increased NO bioavailability can cause the reduced TPR in the GH-supplemented GH-deficient patients.
**eNOS expression and localisation**

Interestingly, in the present study, the level of eNOS protein was unaltered by hypophysectomy in the aorta, caval vein and the heart. Growth hormone deficient patients demonstrate a decreased NO formation in the urine and plasma compared with control patients (4). One explanation for the unaffected vascular eNOS expression in the in vivo studies by Gan et al. (26). Further experiments are needed to elucidate if the effect on eNOS expression in the intima layer of veins (31).

Conclusions

Hypophysectomy per se, is associated with decreased BP and HR. Seven days replacement with GH results in increased eNOS expression in the vasculature, normalisation of HR, but has no effect on the reduced BP. Our results suggest that the induction of eNOS expression

plasma and urine, after chronic GH treatment of GH-deficient patients (4). They suggested that the increase in NO formation was mediated by IGF-I (4). Taken together, our results suggest that the effects on eNOS expression seen in the aorta could be partly mediated by IGF-I and/or by a direct effect of GH on the target cells. We acknowledge that further studies are needed to elucidate if the effect on eNOS expression in the endothelial cells in the aorta is a direct effect mediated by GH and/or IGF-I.

It is also possible that the increased HR (or presumed increased sympathetic activity) seen after both GH and IGF-I treatment could indirectly stimulate eNOS expression in the aorta. An effect of sympathetic activation is less likely since β-receptor blockade appears to stimulate eNOS expression in vitro (25). However, it may be suggested that the increased frequency of cyclic stretching of the aorta, mechanically induced by the increased HR could be responsible for the increased eNOS expression seen in the aorta. This is supported by an ex vivo experiments by Gan et al. (26). Further experiments are needed to elucidate this hypothesis.

Another indirect effect of GH is the enhancement of the conversion of T₄ to tri-iodothyronine (T₃) via up-regulation of deiodinase enzyme (27). The combined treatment of GH and [T₄ + GC] used in the present study may therefore result in increased plasma levels of T₃ (28). In addition T₃ has been shown to induce eNOS expression in the liver of rats (29). Thus, an increased T₃ plasma level may explain the increased eNOS expression in the caval vein after combined treatment with GH and [T₄ + GC] but not after GH alone or after IGF-I. However this possibly confounding effect of T₃ seems to be limited to the caval vein, since changes in aortic eNOS expression was also observed both after GH treatment alone and after GH + [T₄ + GC] treatment.

The immunohistochemical localisation of eNOS showed a specific staining in the aortic endothelial cells of the intima layer in all groups. In contrast, the caval vein demonstrated staining for eNOS in intima, media and adventitia layers. Since von Willenbrand factor has been shown to be specific for endothelial cells, the co-localisation of eNOS and von Willenbrand factor to these cell structures, supports the notion that these cells share some of the features of endothelial cells (30). Thus, it appears that endothelial-like cell types and thereby eNOS localisation is not restricted to the intima layer of veins (31).

Conclusions

Hypophysectomy per se, is associated with decreased BP and HR. Seven days replacement with GH results in increased eNOS expression in the vasculature, normalisation of HR, but has no effect on the reduced BP. Our results suggest that the induction of eNOS expression
seen in the arteries (aorta) after GH treatment could partly be mediated by IGF-I and/or be a direct effect of GH on the vascular cells, whereas in the veins (caval vein) a so far unknown indirect effect of GH may mediate the increased eNOS expression. These findings support the suggested positive role of GH in the regulation of the cardiovascular homeostasis. The observed up-regulation of eNOS, and presumably an increased NO bioavailability, may result in improved endothelial and cardiovascular function.

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References

13 Groesbeck MD & Parlow AF. Highly improved precision of the hypophysectomized female rat body weight gain bioassay for growth hormone by increased frequency of injections, avoidance of antibody formation, and other simple modifications. Endocrinology 1987 120 2582–2590.


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