Quantitative evaluation of CART-containing cells in urinary bladder of rats with renovascular hypertension

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Abstract

Recent biological advances make it possible to discover new peptides associated with hypertension. The cocaine- and amphetamine-regulated transcript (CART) is a known factor in appetite and feeding behaviour. Various lines of evidence suggest that this peptide participates not only in control of feeding behaviour but also in the regulation of the cardiovascular and sympathetic systems and blood pressure. The role of CART in blood pressure regulation led us to undertake a study aimed at analysing quantitative changes in CART-containing cells in urinary bladders (UB) of rats with renovascular hypertension. We used the Goldblatt model of arterial hypertension (two-kidney, one clip) to evaluate quantitative changes. This model provides researchers with a commonly used tool to analyse the renin-angiotensin system of blood pressure control and, eventually, to develop drugs for the treatment of chronic hypertension. The study was performed on sections of urinary bladders of rats after 3, 14, 28, 42 and 91 days from hypertension induction. Immunohistochemical identification of CART cells was performed on paraffin for the UBs of all the study animals. CART was detected in the endocrine cells, especially numerous in the submucosa and muscularis layers, with a few found in the transitional epithelium and only occasionally in serosa. Hypertension significantly increased the number of CART-positive cells in the rat UBs. After 3 and 42 days following the procedure, statistically significantly higher numbers of CART-positive cells were identified in comparison with the control animals. CART appeared in numerous transitional epithelium cells.

As this study provides novel findings, the question appears about the type of connection between hypertension and the functioning and activity of CART in the urinary tract (UT). The study gives rise to the assumption that high blood pressure can be a factor that intensifies CART secretion. In conclusion, the endocrine system of the urinary tract is modified by renovascular hypertension. This may affect the production of hormones and biologically active substances and contribute to the development of possible hypertension complications. In order to fully comprehend the role of the CART peptide in blood pressure regulation, further analyses are necessary.

Introduction

Renovascular hypertension is a common disorder and an illness with an added social dimension. It sporadically occurs as an isolated condition, however more often it occurs in combination with other illnesses, especially metabolic disorders, including disorders of lipid and lipoprotein management, diabetes, obesity, hyperuricemia and changes in the quantitative composition of clotting factors. There are genetic predispositions and disorders of complex mechanisms regulating blood pressure underlying both primary and secondary hypertension. It is postulated that the nervous and endocrine systems, kidneys and local factors regulating the tension of muscles in blood vessels are responsible for regulating the pressure. The results of research in recent years also suggest a considerable role of the endocrine system, including various types of highly specialized, diffusively located receptor-effector cells included in the diffuse neuroendocrine system (DNES). These cells produce active substances that affect the coordination of a wide range of functions of organs and are direct regulators of homeostasis in the body. However, they can simultaneously play roles as mediators in several circulatory disorders, including renovascular hypertension. Among the active substances secreted by DNES cells, there is the cocaine and amphetamine regulated transcript (CART), produced and secreted in the hypothalamus and in other tissues in many vertebrates.4,11 Despite considerable progress made in the area of structure and physiological description of this factor, its role in the pathogenesis of human hypertension is controversial. The development of techniques in molecular biology and histochemistry has enabled research on the distribution and precise tissue localization of the CART peptide in many organs. The principal site of CART synthesis is in areas of the brain, involved in the control of blood pressure.9,11 CART is mainly known for its role in feeding anomalies, which result in hypophagia. This concerns a wide range of biological activities. Among others, CART has neurodegenerative properties after a stroke and is related to blood circulation and renovascular hypertension.10,12-14 Central administration of CART peptide increases systemic blood pressure43 and blocks phenylephrine-induced bradycardia.4 There is only limited information concerning the influence of CART on heart rate, blood pressure, stress and secretion of adrenal hormones. There is both anatomical and physiological evidence available that CART is engaged in hypothalamus - pituitary gland - adrenal gland (HPA) regulation. The HP axis represents one of the main limbs of an adaptive system, which maintains the basal and stress-related homeostasis in vertebrates.112 It is the main component of the neuroendocrine system responsible for the coordination of many systems, including the cardiovascular system.12,14 The fact that CART is found at the level of HPA as well as coexisting with peptides with pressor properties such as leptin, neuropeptide Y (NPY) or calcitonin gene related peptide (CGRP),13 can indicate a potential role of CART in physiological regulation of blood pressure. The presence of CART in the HPA axis is strict-
Description of the experiment

Induction of experimental hypertension was performed according to procedure developed by Goldblatt et al.19 After the rats were anaesthetised by exposure to pentobarbital (40 mg/kg, i.p.), a 3-cm retroperitoneal flank incision was performed in sterile conditions. The left kidney was exposed and the renal artery was carefully dissected free of the renal vein. The renal artery was then partially occluded by placing a standardized silver clamp with an internal diameter of 0.22 mm on the vessel. The wound was closed with a running 3-0 silk suture. The sham-operated rats (n=15) underwent identical surgical procedures, except that a clip was not applied to the renal artery. After the surgery, the rats were kept in separate cages.

After 3, 14, 28, 42 and 91 days from the renal artery clamping procedure, all rats were weighed and the systolic arterial pressure was measured by the tail-cuff method.20 The systolic arterial pressure was calculated by exposure to pentobarbital (40 mg/kg, i.p.), a 3-cm retroperitoneal flank incision was performed in sterile conditions. The left kidney was exposed and the renal artery was carefully dissected free of the renal vein. The renal artery was then partially occluded by placing a standardized silver clamp with an internal diameter of 0.22 mm on the vessel. The wound was closed with a running 3-0 silk suture. The sham-operated rats (n=15) underwent identical surgical procedures, except that a clip was not applied to the renal artery. After the surgery, the rats were kept in separate cages.

After 3, 14, 28, 42 and 91 days following the clamping procedure and induction of renovascular hypertension, all the animals were put into a state of deep anesthesia with pentobarbital (50 mg/kg body weight). After cardiac arrest, fragments of the bladder wall were collected from the animals, always from the same place (body of urinary bladder). Sections of the wall, sampled in the same way as from the experimental group, were collected from rats not undergoing any procedures to provide comparative material.

Tissues were fixed (72 h at room temperature) by immersion in 4% buffered formaldehyde and, following dehydration, were embedded in paraffin wax. The sections were cut 4 μm thick and attached to FLEX IHC microscope slides (K8020, Dako, Glostrup, Denmark).

Immunohistochemical protocol

The identification and visualization of CART in neuroendocrine cells (NE) was performed based on the EnVision method according to Herman and Elfonti using a commercial polyclonal antibody against CART. For antigen retrieval, immunostaining was performed using the below protocol. The sections were deparaffinized in xylene and hydrated in a series of alcohols with decreasing concentration. The sections were subjected to pre-treatment in a pressure chamber heated for 1 min at 21 psi at 125°C using Target Retrieval Solution, pH of 9.0 (S 2367, Dako). After being cooled to room temperature, these sections were incubated with Peroxidase Blocking Reagent (S 2001, Dako) for 10 min to block endogenous activity. The sections with the primary antibody for the CART peptide (rabbit polyclonal CART antiserum, No H-003-61, purchased at the Phoenix Pharmaceuticals, Inc. Mountain View, CA, USA) were diluted (1:10,000) in antibody diluents (S 8989, Dako). The slides were incubated overnight at 4°C in a humidified chamber with the diluted antibody, followed by incubation with a secondary rabbit antibody (conjugated to horseradish peroxidase-labeled polymer) (EnVision+ Kit HRP Rabbit K4011, Dako) for 1 h. The bound antibodies were visualized by 1-min incubation using the 3,3′-diaminobenzidine (DAB) substrate chromogen. All of the sections were finally stained with Vector QB hematoxylin (H-3404; Vector Laboratories, Inc., Burlingame, CA, USA), mounted and evaluated under a light microscope. Appropriate washing with Wash Buffer (S 3006, Dako) was performed between each step.

The specificity test performed for the CART antibody included negative control, where the antibodies were replaced by normal rabbit serum (Vector Laboratories; Burlingame, CA, USA) at the respective dilution. Additionally, positive control was carried out for the specific tissue, as recommended by the producer (for our research we used rat hypothalamus). The stained preparations were analyzed under an Olympus BX 41 microscope. Two sections of each fragment of the urinary bladder from each animal were studied. The cells containing CART were localized and observed in 5 randomly selected fields of vision (0.785 mm²) at a 200× zoom (20× objective and 10× eyepiece) of the analyzed urinary bladder section area.

Statistics

The rats were divided into 2 groups: study and control and each of these 2 into 5 further groups (1-5) according to the duration of the experiment. The duration was measured in days: 1st group, 3 days; 2nd group, 14 days; 3rd group, 28 days; 4th group, 42 days; 5th group, 91 days. After completion of the experiment, the number of CART-immunoreactive (CART-IR) cells in the all layers of the urinary bladder wall was determined for each rat in the particular groups. The parameters were determined at two decimal places accuracy. The determined numbers of endocrine cells (in the individual layers of wall urinary bladder), as well as the duration of illness, were subjected to statistical analysis.

Statistical analysis

All the collected data were analyzed statistically using STATISTICA Version 10.0 (StatSoft Polska Inc., Kraków, Poland). The numbers of NE cells were analyzed using the Mann-Whitney test for independent variables. When testing statistical
hypotheses, P<0.05 was assumed as the level of significance.

Results

No significant differences between the two control groups of rats were observed in the obtained results of the study. Only the results concerning sham-operated animals were taken into account. Blood pressure monitoring showed that, at week 6 of the experiment the blood pressure in the renovascular hypertensive rats stabilized at 162.6±2.19 mmHg, while in control groups remained at 120.2±5.89 mmHg. Hypertensive rats were gaining weight at a similar rate as normotensive animals. Immunohistochemical (IHC) staining enabled the identification of CART-IR cells in the entire urinary bladder wall, both in the normotensive and hypertensive group during all the weeks of induced hypertension. Microscopic observations allowed the location of CART-positive cells, as well as their numbers in relation to the duration of the disorder. Analyzing the location of endocrine cells, in all experimental groups, the most numerous CART-IR cells were found in the urinary bladder muscularis layer and the submucosa (Figure 1A), whereas they were only a few observed in the transitional epithelium and occasionally under urothelium (Figure 1B).

The observed cells were predominantly individually scattered or formed small groups of 2-3 (Figure 1C), sporadically 5-6 cells (Figure 1D).

Quantitative evaluation of CART-IR cells

Statistical analysis confirmed the intuitive view that, in the first stage of the hypertension, the greatest changes included increased activity of NE cells with a significant increase in their mean number in the experimental group (Table 1). An increase of nearly 50% was observed three days after ligation of the left renal artery in the urinary bladder walls of the rats from the experimental group (Figure 2A), in comparison with their counterparts in the control group (Figure 2B).

In the next experimental group, i.e. the two-week one, two weeks after the ligation procedure, a rise in the numbers of NE cells secreting CART was also observed in the control group, though lower when compared with the 1st study group. In the 3rd study group (28 days), endocrine cell density (23.88±6.06) in the animals with renovascular hypertension only slightly differed from the 1st study group. In the analysis, the greatest population of CART-IR cells (in normotensive and hypertensive rats) in the experimental model of unilateral renal artery stenosis was observed after 42 days.

| Time after artery clipping procedure | Group of rats | Number of CART-IR cells |
|--------------------------------------|--------------|-------------------------|
| 3 days                               | Control      | 13.67±2.12              |
|                                      | 2K1C         | 23.63±4.38*             |
| 14 days                              | Control      | 14.67±4.24              |
|                                      | 2K1C         | 22.38±4.06*             |
| 28 days                              | Control      | 15.33±6.71              |
|                                      | 2K1C         | 23.88±6.66*             |
| 42 days                              | Control      | 18.00±1.41              |
|                                      | 2K1C         | 25.50±5.98*             |
| 91 days                              | Control      | 16.00±2.83              |
|                                      | 2K1C         | 21.13±7.06*             |

CART-IR, CART-immunoreactive; 2K1C, Goldblatt’s model of hypertension – two-kidney, one clip model; *P<0.05.

Figure 1. A) CART-IR cells in submucosa and membrane muscularis. B) CART-containing cells in the transitional epithelium and occasionally under urothelium. C) Small groups of 2-3 CART-IR cells in the muscularis layer. D) Group of 5-6 CART-IR cells in the muscularis layer. Scale bars: 20 µm.

Figure 2. A) Three-day form artery clipping procedure; numerous CART-containing cells in the muscularis layer. B) Animals with 3-day, the control group; few NE cells in the muscularis layer. Scale bars: 20 µm.
days of illness in comparison with the other groups included in the experiment. In this group, much larger clusters of cells were also noticed in comparison with the other study groups. The clusters consisted of up to 5-6 cells and were mainly located in the muscularis layer (Figure 3A). Numerous cells were also reported in the urothelium only in this study group. In the mucosa immunoreactive cells were observed in the surface cell layer of transitional epithelium (Figure 3B). At the same time, in the other groups (experimental and control) the cells with positive reaction to the anti-CART antibody in cytoplasm were most often individually scattered or clustered in small groups of 2-3 cells. After 13 weeks from the surgery, the population of CART-IR cells found in the urinary bladder of hypertensive rats significantly decreased in number when compared (21.13±7.06) with the other groups.

Discussion

Renovascular hypertension caused by renal artery stenosis may lead to renal hypoperfusion and development of a sequence of changes in hormone and neuropeptide secretion, which in consequence leads to hemodynamic disorders. Multiple studies have proved that, in patients, renal artery stenosis in early stages of the illness is an incentive to increasing renin production by the ischemic kidney and, as a result, to increased activity of the HPA axis as well as excessive production of angiotensin II.19,22 Among other significant factors in the pathogenesis of hypertension there are active substances correlated with the HPA axis and the sympathetic nervous system. It is assumed that, besides the stimulation of the renal hormonal system, there is an intensified activity of DNES cells, which are diffusely distributed throughout the body and produce pressor substances.23 The group of substances with pressor characteristics, playing a vital role in pathophysiology of hypertension, includes those related to hunger, such as ghrelin, leptin or NPY.24,25

Another substance that has influence on hypertension, and still constitutes a subject of many discussions, is CART. It is principally known for its role in feeding but, as shown in a number of studies, its role is not limited only to hypophagia as it has an impact on many organs and systems.19,24 CART is a peptide involved in a number of physiological processes including modulation of the HPA axis and cardiovascular regulation.15,16,26-30 Its function as a factor in the development of hypertension, despite numerous studies, remains controversial. Arguments in favour of CART influence on the cardiovascular system are provided in studies of Hwang et al.6 and Matsumura et al.19,32 In their experiments, after direct infusion of CART to cerebral ventricles, Matsumura et al.10 noted an increase in mean blood pressure and plasma glucose concentration. On the other hand, liff et al.29 reported that, in isolated cerebral arteries, CART peptide (CARTp) acts directly to produce endothelium-dependent constriction via the endothelin signaling pathway.

Visualization of innervation changes, together with the development of nervous structures secreting CART in the urinary bladder of rats, was performed by Zvarova et al.3 However, the results did not concern normotensive animals. The knowledge relating to the identification and distribution of CART in pathological states caused by hypertension has so far been limited only to specific organs of the alimentary tract, as well as the bladder and ureters.3

Kasacka and Piotrowska postulated the connection of CART with hypertension and used an IHC method in the gastrointestinal tract (Gl tract) of rats in an experimental model of unilateral renal artery stenosis. After 42 days from inducing the condition, the researchers measured the number of NE cells and structures secreting CART in comparison with normotensive animals. The results of the observations were in accordance with the prognosis because the hypertension caused a significant increase in the mean number of CART-IR cells and structures. A close connection between hypertension and the scattered nervous system has also been proved in the region of the lower urinary tract in hypertensive rats. Besides the abovementioned studies, no information has been found on the subject of dynamics of changes in CART-IR cells in the urinary tract of hypertensive animals in relation to the duration of the illness. The connection between the duration of hypertension and the number of DNES cells has been documented only in studies of Kasacka and Arciszewska who analyzed the dynamics of changes in the number of NE cells under the influence of 3-, 14-, 28-, 42-, and 91-day hypertension. The aim of their studies was to examine the distribution, morphology and dynamics of changes of calcitonin gene related peptide-containing cells in the lungs of rats in the two-kidney, one-clip (2K1C) renovascular hypertension model. The authors proved that there is a connection between hypertension, the number of NE cells containing CGRP and the duration of hypertension. Analyzing the results, the greatest intensity of NE cells in the lungs of rats was noticed 14 days after the surgical procedure.

In this paper, the presented data concern the duration of hypertension and the number of CART secreting cells in the urinary bladder of rats in a renovascular hypertension model. After 3 days, the number of NE cells in the urinary bladder walls of the rats that underwent the surgical procedure of inducing renovascular hypertension significantly increased. This pattern was not described by Kasacka and Arciszewska in rat lungs after 2K1C. Three and 28 days after clamping the left renal artery, the number of cells reached comparable levels in the control and study group. However, it was lower in the study group after 3 days since hypertension had been induced. In contrast, Kasacka and Arciszewska observed the greatest number of CGRP-IR cells 14 days after the procedure in the lungs of hypertensive rats. In our observations, the highest parameters were noted only 6 weeks after hypertension was induced. These and earlier promising data let us assume that the studied peptide may play an active role in the development of renovascular hypertension and related disorders.45 Renovascular hypertension caused by renal artery stenosis in original Goldblatt’s experi-
Hypertension, and also modulates fluid balance.36 Renovascular hypertensive rats models, cystometric changes and bladder over-reactivity and urinary incontinence multiple organs.4,6 

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