cAMP/PKA Agonist Restores the Fasting-Induced Down-Regulation of nNOS Expression in the Paraventricular Nucleus

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Gene expression of neuronal nitric oxide synthase (nNOS) changes in the hypothalamic paraventricular nucleus (PVN) depending on feeding conditions, which is decreased during food deprivation and restored by refeeding, and phosphorylated cAMP response element binding protein (pCREB) was suggested to play a role in its regulation. This study was conducted to examine if the fasting-induced down-regulation of the PVN-nNOS expression is restored by activation of cAMP-dependent protein kinase A (cAMP/PKA) pathway. Freely moving rats received intracerebroventricular (icv) injection of cAMP/PKA activator Sp-cAMP (40 nmol) or vehicle (sterilized saline) following 48 h of food deprivation. One hour after drug injections, rats were transcardially perfused with 4% paraformaldehyde, and the PVN tissues were processed for nNOS or pCREB immunohistochemistry. Sp-cAMP significantly increased not only nNOS but also pCREB immunoreactivities in the PVN of food deprived rats. Fasting-induced down-regulation of the PVN-nNOS was restored by 1 h after the icv Sp-cAMP. Results suggest that cAMP/PKA pathway may mediate the regulation of the PVN-nNOS expression depending on different feeding conditions.

Key Words: Food deprivation, Gene expression, Neuronal nitric oxide synthase, Paraventricular nucleus, Rat

INTRODUCTION

Food deprivation decreases mRNA level of neuronal nitric oxide synthase (nNOS), the intracellular intensity of NADPH-diaphorase (NADPH-d) staining, a marker for nNOS activity in the brain, and the apparent number of nNOS immunoreactive cells in the hypothalamic paraventricular nucleus (PVN) [1-5]. Plasma glucocorticoids, which are elevated during food deprivation [3,5-7], appear to mediate the fasting-induced down-regulation of nNOS in the hypothalamic PVN. We have reported that depletion of endogenous glucocorticoids by adrenalectomy abolishes fasting-induced down-regulation of nNOS in the rat PVN, and dexamethasone decreases the PVN-nNOS in both adrenalectomized and intact rats [4,8]. Furthermore, treatment with a glucocorticoids receptor antagonist RU486 during food deprivation blunted the fasting-induced decrease of nNOS in the PVN [3]. These reports together suggested that plasma glucocorticoids may suppress nNOS expression in the PVN during food deprivation, via its receptor-mediated pathway. However, the upstream promoter of nNOS does not carry glucocorticoid response element, but carries cAMP response element [9-11]. In vitro studies have reported that nNOS expression is regulated by a cAMP response element-binding protein (CREB) family transcription factor-dependent mechanism [11], and that phosphorylation of Ser-133 residue in CREB, a critical step in the activation of CREB, is regulated by glucocorticoids [12,13]. We have reported that CREB phosphorylation in the PVN is markedly decreased by 48 h of food deprivation, and this reduction is inhibited by treatment with a glucocorticoid receptor antagonist RU486 [3]. Thus, it is suggested that the plasma glucocorticoids may suppress nNOS expression in the PVN during food deprivation, via down-regulation of CREB phosphorylation.

Not only the fasting-induced decrease of the PVN-nNOS but also of the PVN-pCREB is restored shortly after refeeding [5], and the refeeding-induced increases of the hypothalamic CREB phosphorylation and nNOS expression are blocked by dexamethasone treatments [5,14]. Also, the fast-
ing-induced elevation of the plasma glucocorticoids is restored shortly after refeeding [5,15]. These reports together suggest that refeeding-induced nNOS expression and CREB phosphorylation in the PVN may require glucocorticoid dis-inhibition. However, our previous study demonstrated that glucocorticoid dis-inhibition alone is not sufficient to induce nNOS expression in the PVN [5]. It was reported that food deprivation decreases the hypothalamic cAMP levels and refeeding normalizes it [16]. In this study, we demonstrate that the intracerebroventricular (icv) injection of the membrane-permeable cAMP-dependent protein kinase A (PKA) activator Sp-cAMP (adenosine-3',5'-cyclic monophosphorothioate Sp-isomer) without food supplement rescues the fasting-induced decrease of nNOS expression in the PVN without affecting the plasma corticosterone levels.

**METHODS**

**Animals**

Sprague-Dawley rats were purchased (Samtako Bio, Osan, Korea), and cared in a specific-pathogen-free (SPF) barrier area with constant control of temperature (22±1°C), humidity (55%), and a 12/12 hr light/dark cycle (lights-on at 07:00 AM). Standard laboratory food (Purina Rodent Chow, Purina Co., Seoul, Korea) and membrane filtered purified water were available ad libitum. Animals were cared according to the Guideline for Animal Experiments, 2000, edited by the Korean Academy of Medical Sciences, which is consistent with the NIH Guidelines for the Care and Use of Laboratory Animals, revised 1996. All animal experiments were approved by the Committee for the Care and Use of Laboratory Animals at Seoul National University.

**Intracerebroventricular (icv) cannulation**

Under chloral hydrate (153 mg/kg) and pentobarbital (35 mg/kg) anesthesia, rats (280−300 g) were stereotaxically implanted with a 22-gauge, stainless steel guide cannula (Plastics One, Roanoke, VA) aimed towards the lateral ventricle (1.2 mm caudal to bregma, 1.5 mm lateral to the midline, and 4 mm below the skull surface). Guide cannulae were held in place with dental acrylic bonded to stainless steel screws anchored to the skull. An obdurator was inserted into each guide cannula and remained in place except during injections when it was removed and replaced with an injector that extended 1.0 mm beyond the tip of the guide cannula. After 1 week of post-operational recovery, patency and placement of the cannula was verified by injection of 100 ng human angiotensin II (Sigma Chemical Co., St Louis, MO, USA) dissolved in 5 μl of 0.15 M NaCl; rats with cannulas projecting into the lateral ventricle responded to the angiotensin injection by vigorously licking the water bottle within 2 min [17-19], while rats that failed to drink were dropped from the study. Cannula placements were also verified postmortem by sectioning through the brain.

**Drug treatments**

Rats verified for the cannula placement were divided into four different groups (n=6 in each group, total 24 rats); i.e., FC/vehicle (saline icv with ad libitum access to food), FD/vehicle (saline icv following food deprivation), FD/Sp-cAMP (Sp-cAMP icv following food deprivation) and FD/Rp-cAMP (Rp-cAMP icv following food deprivation). Rats in FD/vehicle, FD/Sp-cAMP or FD/Rp-cAMP groups underwent a 48 h of food deprivation beginning at 09:00 h, and then received an icv injection of Sp-cAMP (Sp-8-bromo-cAMP, Sigma Chemical Co., St Louis, MO, USA; 5 μl of 40 nmol in sterilized saline), Rp-cAMP (Rp-8-bromo-cAMP, Sigma Chemical Co., St Louis, MO, USA; 5 μl of 40 nmol in sterilized saline) or sterilized saline (5 μl of 0.15 M NaCl). Sp-cAMP, Rp-cAMP or saline was delivered through the icv guide cannulas over 30 s with a handheld 50 μl syringe (Hamilton Co., Reno, NV, USA). The injector was left in place for 30 s after solution delivery. Free-fed control group (FC/vehicle) received an icv injection of sterilized saline (5 μl of 0.15 M NaCl).

**Immunohistochemistry**

One hour after the drug injections, rats were anesthetized with over doses of sodium pentobarbital (Hallym Pharmaceutical Co., Seoul, Korea), and transcardiac perfusions were performed first with heparinized isotonic saline then with ice-cold 4% paraformaldehyde (Sigma Co., MO, USA) in 0.1 M sodium phosphate buffer. Brains were immediately dissected out, blocked, post-fixed for 2 h, and transferred into 30% sucrose (Sigma Co., MO, USA) for cryoprotection. Forty-micron coronal sections were cut on a freezing, sliding microtome (HM440E, Microm Co., Germany). Alternate sections were collected throughout the rostrocaudal extent of the hypothalamic PVN (between bregma - 1.3 mm and - 2.1 mm) [20]. Immunohistochemistry was performed with standard DAB reaction using commercial ABC kit (Vectastain Elite Kit, Vector Laboratories, CA, USA) as previously described [21]. Polyclonal rabbit anti-nNOS, which reacts with the 160 kDa nNOS protein and does not exhibit any cross-reactivity with the related eNOS or iNOS proteins (1: 2,000 dilution, Zymed Laboratories, CA, USA), and antigen retrieval was performed by microwave heating in citrate buffer (pH 6.0). Sections were then incubated with pCREB (1: 2,000, Upstate biotechnology, NY, USA), antibodies were used as primary antibodies, and biotinylated anti-rabbit IgG (1: 200 dilution, Vector Laboratories, CA, USA) as secondary. Immunostained sections were mounted in an anatomical order onto gelatin-coated slides from 0.05 M phosphate buffer, air-dried, dehydrated through a graded ethanol to xylene, and cover-slipped.

**Plasma corticosterone assay**

Cardiac blood was rapidly collected in the heparinized tube immediately after exposing the heart for perfusion, and centrifuged at 2,000 rpm for 20 min. The plasma was transferred into new tubes, frozen in liquid nitrogen, and stored at −80°C until used for the assay. Plasma corticosterone levels were determined by radioimmunoassay using 125I-labelled Coat-A-Count kit (DPC, CA, USA). All rats were sacrificed between 10:00 h and 12:00 h to minimize diurnal variation in the plasma levels of corticosterone [22].

**Statistical analysis**

The number of nNOS immunopositive cells was blind-counted by hand, pCREB immunopositive nuclei auto-counted, after digitizing 720×540 micron images of three sections from the PVN (closest sections to bregma - 1.88 mm) from each brain using an Olympus BX-50 microscope.
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Fig. 1. nNOS immunoreactivity and number of nNOS-ir neurons in the parvocellular and the magnocellular PVN. Rats were sacrificed for nNOS immunohistochemistry at 1 h after an intracerebroventricular (icv) injection of Sp-cAMP or saline vehicle. FC/veh, free-fed control rats received icv administration of 5 μl sterilized saline; FD/veh, 48 h food deprivation and received icv administration of 5 μl sterilized saline; FD/Sp-cAMP, 48 h food deprivation and received icv administration of 5 μl of 40 nmol Sp-cAMP in sterilized saline; FD/Rp-cAMP, 48 h food deprivation and received icv administration of 5 μl of 40 nmol Rp-cAMP in sterilized saline; mP, medial parvocellular subdivision; pM, posterior magnocellular subdivision; PVN, paraventricular nucleus; 3v, third ventricle; *p < 0.05, **p < 0.01 vs. FC/veh; Values are presented as means±S.E.

Fig. 2. pCREB immunoreactivity and number of pCREB-ir nuclei in the parvocellular and the magnocellular PVN. Rats were food deprived for 48 h and then received icv administration of 5 μl sterilized saline (vehicle) or 40 nmol Sp-cAMP in sterilized saline, and sacrificed 1 h after the icv injections for pCREB immunohistochemistry. mP, medial parvocellular subdivision; pM, posterior magnocellular subdivision; PVN, paraventricular nucleus; 3v, third ventricle; *p < 0.05 vs. vehicle controls; Values are presented as means±S.E.

Fig. 3. Plasma corticosterone levels at 1 h after an icv administration of Sp-cAMP or saline vehicle. FC/veh, free-fed control rats received icv administration of 5 μl sterilized saline; FD/veh, 48 h food deprivation and received icv administration of 5 μl of 40 nmol Sp-cAMP in sterilized saline; FD/Sp-cAMP, 48 h food deprivation and received icv administration of 5 μl of 40 nmol Sp-cAMP in sterilized saline; *p < 0.05 vs. FC/veh, Values are presented as means±S.E.

Results

Food deprivation significantly reduced the number of nNOS immuno-positive (ir) neurons in both the medial parvocellular (mP) and the posterior magnocellular (pM) PVN (Fig. 1). One hour after an icv Sp-cAMP, nNOS-ir was significantly increased in the parvocellular PVN of food deprived rats (p < 0.05, FD/veh vs. FD/Sp-cAMP); however, this increase was not observed in the magnocellular PVN. nNOS-ir in the PVN of food deprived rats that received an icv Rp-cAMP, cAMP antagonist, did not differ from the food deprived rats that received icv saline.

pCREB-ir in the parvocellular PVN of food deprived rats that received icv Sp-cAMP was increased significantly (p < 0.05) compared with vehicle injected controls (Fig. 2). Sp-cAMP did not increase pCREB-ir in the magnocellular PVN of food deprived rats.

Cardiac blood was collected at sacrifice and the plasma levels of corticosterone were analyzed by radioimmunoassay (Fig. 3). Food deprivation significantly increased the plasma levels of corticosterone (p < 0.05, FC/veh vs. FD/veh), and Sp-cAMP administration did not alter the plasma corticosterone levels of food deprived rats.


DISCUSSION

We have previously demonstrated that nNOS gene expression in the hypothalamic PVN changes depending on feeding conditions; i.e., it is decreased during food deprivation and restored following refeeding, and that pCREB may be involved in the PVN-nNOS expression [3,5]. The upstream promoter of nNOS gene contains cAMP response element (CRE) [9]. In vitro studies have reported that nNOS expression is regulated by calcium influx through a CREB family transcription factor-dependent mechanism [11], and that cAMP markedly induces nNOS expression in human A673 neuroepithelial cells expressing CREB, and the induction is enhanced in A673 cells transfected with CREB [23]. In vivo study showed that the hypothalamic cAMP levels are decreased during food deprivation and restored by food consumption [16]. In this study, icv administration of cAMP-dependent PKA activator Sp-cAMP increased not only pCREB levels but also nNOS expression in the PVN of food deprived rats, and cAMP antagonist/PKA inhibitor Rp-cAMP [16] did not affect the fasting-induced down-regulation of the PVN-nNOS. Together, it is suggested that the regulation of the PVN-nNOS expression depending on feeding conditions may be mediated by cAMP/PKA pathway. Previous studies have suggested that the plasma glucocorticoids may suppress nNOS expression in the PVN during food deprivation, via down-regulation of CREB phosphorylation [3,4,8], and that glucocorticoids dis-inhibition is required, although it is not sufficient, to restore the fasting-induced down-regulation of the PVN-nNOS and -pCREB [5,14].

However, in this study, Sp-cAMP administration increased both nNOS and pCREB in the PVN of food deprived rats, despite the fasting-induced elevation of the plasma corticosterone was still persisted. In vitro studies have reported that glucocorticoids regulate phosphorylation of Ser133 residue in CREB, a critical step in activation of CREB [12,13], and interfere with the cAMP-induced DNA binding of CREB [24]. Underlying mechanism by which Sp-cAMP increased pCREB and nNOS in the PVN of food deprived rats despite of the glucocorticoids inhibition is currently not clear. It should be noticed that cAMP agonist used in this study Sp-cAMP is a more potent PKA activator with higher affinity to the regulatory subunit of PKA and more resistant to phosphodiesterase than endogenous cAMP [16,25], which might have allowed Sp-cAMP to overcome the glucocorticoids inhibition.

Our previous study suggested that refeeding-induced nNOS expression in the PVN is a nutrient directed event; i.e., refeeding with non-caloric palatable food mash failed to restore the fasting-induced decreases not only of the PVN-nNOS but also of -pCREB [5]. Many studies have examined the effects of various nutrients on NOS expression, however, those studies have been mostly focused on endothelial or inducible NOS, but not nNOS [26]. In vitro studies have demonstrated that glucose, the major substrate for cerebral energy metabolism, increases the intracellular cAMP content [27], and activates cAMP/PKA pathway [28]. Also, an intragastric infusion of glucose induced c-Fos expression, conventional marker for neuronal activation, in the PVN of food deprived rats [29]. Together with the current results, it is suggested that postprandial glucose may induce NOS expression in the PVN at refeeding following food deprivation, by activating the cAMP/PKA pathway.

In this study, food deprivation down-regulated nNOS expression both in the parvocellular and the magnocellular di-visions of PVN; however, Sp-cAMP restored NOS expression only in the parvocellular PVN, but not in the magnocellular PVN. Furthermore, Sp-cAMP following food deprivation increased pCREB levels only in the parvocellular PVN, but not in the magnocellular PVN. Our previous study showed that CREB phosphorylation is associated with nNOS down-regulation in the parvocellular PVN, but not in the magnocellular PVN, during food deprivation. That is, the fasting-induced nNOS down-regulation was observed not only in the parvocellular PVN but also in the magnocellular PVN; however, food deprivation reduced pCREB only in the parvocellular PVN [3]. Thus, it is concluded that NOS expression in the magnocellular PVN, not likely in the parvocellular PVN, may not be mediated by cAMP/PKA pathway.

Nitric oxide (NO) is an important biological messenger that acts in a vast array of physiological processes [30]. In the nervous system, neuronal nitric oxide synthase (nNOS) accounts for the majority of the physiologic actions of NO [31,32]. Many investigations have shown that NOS expression is dynamically regulated by both physiological and pathophysiological stimuli [31,33-35]. Feeding-related changes of nNOS expression in the PVN may be a part of the regulatory pathway for autonomic changes associated with feeding status, such as heart rate and blood pressure [36]. The PVN nitric oxide has been suggested to be involved in the regulation of autonomic functions [37,38], and reported to modulate vasopressin release [39]. Vasopressin is known to be involved in the control of blood pressure and heart rate as well as in the regulation of fluid and electrolyte balance [40]. Food deprivation decreases the hypothalamic contents of vasopressin [41] and refeeding activates vasopressin neurons in the PVN [15]. Overall, it is postulated that feeding-related changes in NOS expression in the PVN may play a role in the regulatory control for blood pressure and heart rate, at least partly modulating the activity of vasopressin neurons.

ACKNOWLEDGEMENTS

This study was supported by grants from the Brain Research Center of the 21st Century Frontier Research Program (2012K001115) and the National Research Foundation (2010-0003642) funded by the Korea Government (Ministry of Education, Science and Technology).

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