Involvement of brain-derived neurotrophic factor in the regulation of hypothalamic somatostatin *in vivo*

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Abstract

Brain-derived neurotrophic factor (BDNF) has been extensively studied in the central nervous system as a survival and differentiation factor and in plasticity processes. *In vitro*, BDNF has been shown to stimulate cellular differentiation and neurohormones synthesis and release. We demonstrated that BDNF is a potent and specific stimulatory agent of somatostatin (SRIH) synthesis in primary cultures of hypothalamic neurons. However, less information is available about its function on SRIH neurons *in vivo*. In the present study, we examined the effect of *in vivo* intracerebroventricular BDNF administration in adult non-anesthetized male rats. Two distinct experimental approaches were used: acute intracerebroventricular injection and long-term (14 days) continuous infusion (Alzet micro-pumps). We demonstrate that single intracerebroventricular BDNF injections (5 µg/rat)

induce an early (60 and 180 min) decrease in the SRIH mRNA signal in the hypothalamic periventricular nucleus (PeVN) accompanied by a decrease of the hypothalamic SRIH content. 48 h after the acute injection, SRIH mRNA levels and peptide content strongly and significantly increased. After continuous intracerebroventricular BDNF administration (12 µg/day for 14 days), a significant increase in the SRIH hypothalamic content was observed. Nevertheless, the increase in peptide content was not correlated with a similar increase in the PeVN messenger level. These findings show the involvement of BDNF in the *in vivo* regulation of somatostatinergic neurons in adult rats, which clearly differs according to the BDNF administration mode.

Journal of Endocrinology (2006) **188**, 425–433

Introduction

The brain-derived neurotrophic factor (BDNF) is a member of the neurotrophin family having the nerve growth factor as prototype. The main biological effects of BDNF are mediated by specific tyrosine kinase (TrkB) receptors. In adult rats, BDNF is involved in a variety of forms of synaptic plasticity in the central nervous system (CNS). Among these actions, particular interest has been focused on its stimulatory effect on peptide biosynthesis in the CNS (Tapia-Arancibia *et al.* 2004). BDNF has thus been shown to increase mRNA and peptide levels of neuropeptide Y and somatostatin (SRIH) in cultured cortical (Nawa *et al.* 1993) and striatal (Mizuno *et al.* 1994) neurons. In BDNF-knockout mice, the density of SRIH cells is drastically reduced in hippocampus and cortex (Große *et al.* 2005). In hypothalamic cultures, BDNF has been reported to have a stimulatory effect on SRIH content and mRNA expression (Rage *et al.* 1999) and on differentiation (Loudes *et al.* 1999).

In addition, the fact that both SRIH (Benyassi *et al.* 1993a,b) and BDNF (Rage *et al.* 2002, Givalois *et al.* 2004) are molecules involved in the stress response and in food intake regulation (Feifel & Vaccarino 1994, Pellemounter *et al.* 1995) is an important clue in suggesting a functional relationship between these signals. Some of these changes occur in the periventricular nucleus (PeVN), which is a hypothalamic region where BDNF and TrkB mRNAs (Marmigère *et al.* 1998) and most neuronal perikarya secreting SRIH are localized (Makara *et al.* 1983).

This data strongly suggests that BDNF might be a signal involved in the control of SRIH *in vivo*. We assessed this possibility by administering a single BDNF i.c.v. injection in adult rats and analyzing SRIH mRNA levels in the PeVN, the main source of SRIH in the hypothalamus (Brownstein *et al.* 1975). To compare this treatment with the effects obtained by long-term BDNF infusion, similar studies were also carried out on animals bearing osmotic pumps (Alzet) chronically implanted in the lateral ventricle, which allows delivery of a sustained and constant amount of BDNF into the brain.

Materials and Methods

Animals

Adult male Sprague–Dawley rats (Depré, St Doulchard, France) weighing 260–280 g were housed for 1 week under constant temperature ($21 \pm 1^\circ\text{C}$) and lighting (light on from 07:00 to 19:00) regimens. Food pellets and water were freely available. Procedures involving animals and their care were conducted in compliance with French laws on laboratory animals that, in turn, are in compliance with international laws and policies (EEC Council Directive 86/609, OJ L 358, 1, 12 December 1987, NIH Guide for the Care and Use of Laboratory Animals, NIH Publication No. 85–23 1985). All protocols were approved by the Animal Welfare Committee at the University of Montpellier II, which aims to minimize the number of animals used, and potential pain and distress.

Experimental procedures

As repeatedly reported (Lapchak & Hefti 1992, Pellemounter *et al.* 1995, Siuciak *et al.* 1996), we infused BDNF in the lateral ventricle to enable BDNF access to a wide range of periventricular regions, including direct albeit limited access to forebrain areas such as the hippocampus and hypothalamus (Yan *et al.* 1994, Anderson *et al.* 1995). This mode of administration produces minimal lesion damage. At the end of each experiment, the correct cannula implantation and tissue integrity was checked, and rats with the wrong cannula localization or presenting lesions were not included in this study. The choice of sodium chloride as vehicle was based on a study of Callahan *et al.* (2001) showing that the utilization of NaCl enhanced the shelf-life and conformational stability of BDNF.

Surgery

Single i.c.v. injection As previously reported for i.c.v. injection studies (Givalois *et al.* 2004), 7 days before the experiments, the animals were deeply anesthetized with an intramuscular injection of 0.2 ml of a mixture of ketamine hydrochloride (80 mg/kg body weight (b.w.)) and xylazine (10 mg/kg b.w.). They were then stereotactically implanted with a permanent stainless-steel cannula (PlasticOne Products Co., Roanoke, VA, USA) into the left lateral ventricle of the brain at coordinates (AP: –1 mm, L: ± 1.5 mm and DV: –3.5 mm) according to Paxinos and Watson (1997). The rats were caged separately, handled every day before the experiments and had recovered their preoperative body weight prior to i.c.v. injection. On the day of the experiment, freely moving animals were divided into three experimental groups, i.e. control rats (with no cannula), sham rats that received a single i.c.v. injection of saline (5 μl of 0.9% NaCl), and treated rats that received a single i.c.v. injection of

recombinant human BDNF (5 $\mu\text{g}/\text{rat}$ in 5 μl of NaCl 0.9%; kindly provided by Regeneron Pharmaceuticals Inc., Tarrytown, NY, USA). The BDNF dose was selected on the basis of previous experiments, wherein adequate diffusion of this molecule from lateral ventricle to the periventricular areas was considered (Yan *et al.* 1994). The effects of i.c.v. injection on the different parameters studied were determined before (t_0) and 30, 60, 180, 300 min and 48 h after injection.

Continuous i.c.v. injection The animals were also divided in three groups: control, sham and treated. The control group was composed of intact untreated rats, whereas sham and treated rats had an osmotic pump (Alzet model 2002, 200 μl ; 0.5 $\mu\text{l}/\text{h}$; Charles River, France) implanted in the lateral ventricle, as previously reported (Pellemounter *et al.* 1995). Before surgery, each rat was intra-muscularly anaesthetized with a ketamine (80 mg/kg b.w.) and xylazine (10 mg/kg b.w.) mixture. For the osmotic pump implantation, each animal was placed on the stereotaxic apparatus to implant a cannula into the left lateral ventricle, as detailed above. The cannula was sealed with dental cement and connected to an Alzet pump by medical grade vinyl tubing. The pump was placed into a subcutaneous pocket in the dorsal region. For the sham group, the pumps were filled with vehicle (NaCl 0.9%) and for the treated group with a BDNF solution (1 $\mu\text{g}/\mu\text{l}$; kindly provided by Regeneron Pharmaceuticals Inc.). Animals implanted with osmotic pump filled with BDNF received 12 $\mu\text{g}/\text{day}$ of BDNF for 14 days. The pumps were filled the day before surgery was performed. The pumps and the tubing were incubated at 37°C overnight in a sterile saline solution to prime them. The experiment continued for 14 days after pump implantation. The BDNF dose was selected on the basis of previous experiments (Siuciak *et al.* 1996) and this protocol was validated by the loss of body weight observed in treated animals (Lapchak & Hefti 1992, Pellemounter *et al.* 1995, Siuciak *et al.* 1996), which indicates a BDNF access to the hypothalamic areas.

Preparation of digoxigenin-labeled oligonucleotide probe

As previously reported (Arancibia *et al.* 2001), a 45-mer oligonucleotide antisense probe for somatostatin-14 (5' to 3' sequence: CCAGAAGAAAGTTCTTGCAGCCAGCTTTGCGTTCCCGGGGTGT) (Genosys, Cambridge, UK) was end-labeled with digoxigenin-11-dideoxyuridine-5'-triphosphate. Negative controls were carried out by omitting the labeled probe from the hybridization buffer, or by incubating the sections with a 45-base oligonucleotide sense probe based on the somatostatin-14 sequence. Labeling was carried out according to the same protocol as described for the antisense probe, same concentration as the antisense probe were used.

In situ hybridization

On the day of the experiments, the animals were deeply anesthetized with an intramuscular injection of 0.2 ml of a mixture of ketamine hydrochloride (80 mg/kg b.w.) and xylazine (10 mg/kg b.w.) and then rapidly perfused transcardially with 4% paraformaldehyde in 0.2 M phosphate buffer. Brains were removed and postfixed in the same fixative for 4 h at 4 °C, and then placed in 15% sucrose overnight at 4 °C. Thereafter, the tissues were quickly frozen in isopentane chilled in liquid nitrogen. The frozen brains were mounted on a cryostat (Leica, Rueil-Malmaison, France) and serially cut into 10 µm coronal sections. The hypothalamic sections were mounted on Superfrost-Plus glass slides (Menzel-Glaser, Labonord, Templemars, France) and kept at -80 °C until use.

In situ hybridization analysis was carried out as previously described (Arancibia *et al.* 2001). Briefly, the paraformaldehyde-fixed sections were washed for 5 min in PBS before incubation at room temperature for 10 min with 10 mg/ml of proteinase K. The sections were then acetylated with 0.25% acetic anhydride for 10 min. They were then dehydrated in an ascending series of ethanol concentrations and air-dried. The sections were hybridized overnight in a humid chamber at 37 °C with 3 ng of SRIH-labeled probe in 30 µl of hybridization mixture. After hybridization, the sections were then rinsed briefly and incubated at room temperature with 2% (w/v) blocking reagent: 0.1% diethyl pyrocarbonate for 30 min to block nonspecific antibody reactions. Slides were then transferred to a humid chamber and incubated with a 1:500 dilution of alkaline phosphatase-conjugated anti-digoxigenin antibody (Roche) containing 2% (w/v) non-fat dry milk for 2 h. This was followed by two 15-min washes before transfer to a color developing solution, containing nitroblue tetrazolium salt and 5-bromo-4-chloro-3-indolyl phosphate. The color reaction was stopped after 10 h by washing in 200 ml of 1 × Tris-EDTA buffer, pH 8, for 30 min. Excess Tris buffer was wiped off, and the sections were coverslipped with Entellan (Merck).

Image analysis

We took advantage of the non-isotopic *in situ* hybridization technique, which is a valuable method to quantify the results as initially reported from Kiyama and Emson (1990). Thus, the histological sections of the PeVN region were analyzed by counting all labeled cells/section and by measuring the optical density (OD) variations using a Sony CCD XC-77 video camera with high-resolution (570 (H) 3 485 (V) TV lines) coupled to a Macintosh computer (Power PC G3) and NIH-Image software (version 1.63 non-FPU, W. Rasband, NIH, Bethesda, MD) (Arancibia *et al.* 2001, Givalois *et al.* 2004). For each animal, an OD and a number of positive cells/section

from four to six sections measured on each side of the PeVN were determined to calculate a mean per animal. The OD of each specific region was corrected for the average background signal determined by sampling cells immediately outside the cell groups of interest. The experiments were performed twice independently and all sections from control, sham and treated animals were hybridized at the same time to avoid intrinsic variations between different *in situ* hybridizations.

Hypothalamic SRIH content

After decapitation, hypothalami were dissected and immediately frozen on dry ice and stored at -20 °C until the SRIH assay. SRIH hypothalamic contents were measured with a conventional RIA. The SRIH RIA kit (Phoenix Pharmaceuticals Inc., Belmont, USA) was used according to the manufacturer's protocol. The assay sensitivity was 2 pg/tube. SRIH content was expressed as pg/hypothalamus. The intra- and inter-assay coefficients of variation were 4% and 7%, respectively.

Statistical analysis

Data was presented as mean ± S.E.M. Mean and S.E.M. were calculated from 5–6 animals per group for SRIH assay and for the *in situ* hybridization analysis. The different groups were compared by performing an ANOVA (Statview 4.5) followed by a Fisher's PLSD test, as previously reported (Arancibia *et al.* 2001, Givalois *et al.* 2004). $P < 0.05$ was considered significant.

Results

Effect of single i.c.v. BDNF injection on SRIH mRNA and SRIH hypothalamic content

Figure 1 shows time-course variations in SRIH mRNA signals in the PeVN in controls and 60, 180, 300 min and 48 h after BDNF injection.

The SRIH mRNA signal was evaluated in the PeVN, the hypothalamic region containing most of somatostatin-ergic neuron perikarya. Quantifications of messenger variations are represented in Fig. 2A and 2B. BDNF injection rapidly and significantly ($F_{4,18} = 146.7$; $P < 0.0001$) decreased SRIH mRNA signal (OD) 60, 180 and 300 min after BDNF injection (-23% at 60 min, $P < 0.01$ versus control (t_0); -17% at 180 min, $P < 0.01$ versus control (t_0) and -20% at 300 min, $P < 0.01$ versus control (t_0)). By contrast, at 48 h SRIH messenger levels were significantly increased by 20% ($P < 0.01$ versus control rat (t_0)) (Fig. 2A). The number of positive cells expressing SRIH mRNA in the PeVN was significantly decreased ($F_{4,17} = 23.02$; $P < 0.0001$) 60 and 180 min after BDNF injection (-19% at 60 min, $P < 0.01$ versus control rat (t_0))

Figure 1 Effects of single i.c.v. BDNF injection (5 µg/rat) on SRIH mRNA expression in the PeVN, detected by *in situ* hybridization. Figures show representative photomicrographs of histological sections (10 µm) hybridized with digoxigenin-labeled SRIH oligonucleotide probe in control adult male rats (a and b), 60 (c and d), 180 (e and f), 300 min (g and h) and 48 h (i and j) after a single i.c.v. injection of BDNF (5 µg/rat). Photomicrographs a, c, e, g and i: scale bar=400 µm and photomicrographs b, d, f, h and j: scale bar=100 µm. 3v: third ventricle.

and -12% at 180 min, $P < 0.05$ versus control rat (t_0)). At 300 min, the number of positive cells/section was comparable to the control value and was increased by 15%, 48 h after the treatment ($P < 0.05$ versus control rat (t_0))) (Fig. 2B). Figure 2C shows time-course variations of hypothalamic SRIH contents in control animals (t_0) and 30, 60, 180 min and 48 h after a single BDNF injection. The SRIH content was progressively and significantly decreased ($F_{3,27} = 3.77$; $P < 0.022$) 60 and 180 min after BDNF injections (18.7 ± 2.7 ng/hypothalamus at 60 min, $P < 0.01$ and 23.6 ± 1.9 ng/hypothalamus at 180 min, $P < 0.05$ versus 37.9 ng/hypothalamus in control rats (t_0)). By contrast, at 48 h the SRIH content was significantly increased ($F_{1,14} = 18.39$; $P < 0.0008$) by 54% (56.9 ± 3.9 versus 37.9 ± 3.9 in control rats (t_0))) (Fig. 2C).

In situ hybridization of SRIH mRNA as well as SRIH hypothalamic content variations were performed in control, sham and BDNF-treated rats. No difference was observed between control and sham groups. Hybridizations with the sense probes did not reveal any positive signal.

Effect of continuous i.c.v. BDNF administration on SRIH mRNA and SRIH hypothalamic content

Figure 3 shows that, as expected, long-term BDNF treatment (i.c.v. infusion of BDNF, 12 µg/days for 14 days) has decreased body weight compared with control and

sham rats ($F_{2,10} = 17.69$; $P < 0.0005$) indicating BDNF access to hypothalamic areas as already established (Lapchak & Hefti 1992). This decrease was significant from 7 days of treatment (341 ± 15 versus 398 ± 14 g; $P < 0.05$ versus control rats) and at the 14th day, BDNF-treated animals weighed 29% less than control rats (301 ± 21 versus 424 ± 15 g; $P < 0.01$ versus control rats). Figure 4 shows variations in SRIH mRNA signals in the PeVN of controls and treated animals. Quantification of SRIH mRNA levels (Fig. 5A) and the number of positive cells/section (Fig. 5B) showed this treatment decreased SRIH mRNA steady state levels ($F_{1,8} = 54.6$; $P < 0.0001$) by 14% ($P < 0.01$ versus control rats; Fig. 5A) and the number of SRIH mRNA labeled neurons/section ($F_{1,8} = 15.2$; $P < 0.0046$) by 24% ($P < 0.01$ versus control rats; Fig. 5B). In contrast, the SRIH content was significantly increased ($F_{2,10} = 4.47$; $P < 0.041$) by 47% after BDNF treatment (63.8 ± 6.6 ng/hypothalamus versus 43.5 ± 6.3 ng/hypothalamus in control rats; $P < 0.05$) (Fig. 5C).

No differences in mRNA level and in SRIH content were observed between control and sham groups, and hybridizations with the sense probes did not reveal any positive signal.

Discussion

The major finding of this work is the evidence of an *in vivo* regulation of hypothalamic SRIH neurons after

Figure 2 Effects of single i.c.v. BDNF injection (5 µg/rat) on hypothalamic SRIH. (A) and (B) represent the optical density (OD) variations of the SRIH mRNA signal and the positive cells number/section respectively, for the SRIH mRNA signal in the PeVN in control adult male rats (t_0), 60, 180, 300 min and 48 h after the i.c.v. injection of BDNF. The results are expressed as means $\pm$ S.E.M. of the percent of control (considered as 100%), with $n=5-6$ rats per experimental group. * $P<0.05$ and ** $P<0.01$ vs. control group. (C) represents the time-course profiles of hypothalamic SRIH content in control rats (t_0) and 30, 60, 180 min and 48 h after a single i.c.v. injection of BDNF. The results are expressed as means $\pm$ S.E.M. with $n=5-6$ rats per experimental group. * $P<0.05$ and ** $P<0.01$ versus basal value at t_0 .

Figure 3 Time-course variation of body weight (g) during 14 days in control, saline (NaCl 0.9%) or BDNF (12 µg/day per rat) treated rats. The rats were continuously infused using osmotic micro-pumps (Alzet) as described in the experimental procedures. The results are expressed as means $\pm$ S.E.M. with $n=5-8$ rats per experimental group. * $P<0.05$ and ** $P<0.01$ versus control group.

acute or chronic BDNF administration. After a single BDNF injection, we observed a rapid decrease in SRIH mRNA levels and SRIH mRNA positive cells in the PeVN as well as a decrease in SRIH hypothalamic content. In contrast, 48 h later BDNF exerted a stimulatory effect on all of these parameters. These late effects are in line with our previous data showing a BDNF stimulatory effect on SRIH synthesis in cultured hypothalamic neurons after a 48 h treatment (Rage *et al.* 1999).

Among all of the SRIH synthesizing hypothalamic nuclei, the PeVN largely contains the highest number of hypothalamic SRIH perikarya (Alpert *et al.* 1976) and the highest amounts of brain SRIH are found in the median eminence, which result from the PeVN synthesizing activity (Brownstein *et al.* 1975). Therefore, and in accordance with other authors (Critchlow *et al.* 1981, Kiyama & Emson 1990), it is reasonable to consider that a variation in SRIH hypothalamus content is a good functional reflect of SRIH synthesizing activity occurring in PeVN. Thus, a decreased SRIH content could be associated with a rapid stimulatory effect of BDNF on SRIH release, in accordance with our previous *in vitro* data (Marmigère *et al.* 2001).

Considering the high BDNF levels in the adult hypothalamus (Katoh-Semba *et al.* 1997, Rage *et al.* 2002) and the high expression of its mRNA and of its receptor in the PeVN (Marmigère *et al.* 1998), our present results provide solid evidence that BDNF has a physiological role in this structure. Moreover, we have reported high levels

Figure 4 Effects of continuous BDNF administration (12 μ g/day per rat for 14 days) on hypothalamic SRIH mRNA expression in the PeVN, detected by *in situ* hybridization. Hypothalamic sections (10 μ m) were hybridized with digoxigenin-labeled SRIH oligonucleotide probe in control adult male rats (A and B) and in rats 14 days after continuous i.c.v. BDNF administration (C and D). Figures show representative photomicrographs of the PeVN sections hybridized with the SRIH probe. Photomicrographs A and C: Scale bar=300 μ m and photomicrographs B and D: Scale bar=80 μ m. 3v: third ventricle.

of TrkB mRNA in the hypothalamus (Tapia-Arancibia *et al.* 2001) and demonstrated, by double immunocytochemical staining experiments, that TrkB receptors are located on somatostatinergic neurons (Rage *et al.* 1999). As a whole, we can infer that the effect reported here is probably direct, exerted by BDNF on these neurons in an autocrine or paracrine fashion. However, we cannot rule out the potential role of other endogenous neurotransmitters released after BDNF administration.

Nawa *et al.* (1994) reported that *in vivo* administration of BDNF in newborn rat brain increased the contents of several peptides including SRIH in brain cortex, striatum and hippocampus. These authors showed that intraventricular injection of BDNF linearly increased SRIH mRNA expression in the anterior neocortex, reaching maximal levels at 48 h. Nevertheless, these authors failed to detect any marked influence of BDNF on SRIH or neuropeptide Y in the hypothalamus. The differences relative to our results could be explained by the fact that at neonatal stages, endogenous neurotrophin levels are sufficient to ensure peptidergic expression and exogenous administration of BDNF would therefore not further alter the peptide phenotype.

The second paradigm used here, i.e. long-term i.c.v. BDNF administration reproduced the stimulatory effect of BDNF on SRIH content described here (48 h after the single BDNF injection) and those described in previous *in vitro* studies (Rage *et al.* 1999, Loudes *et al.* 2000). Using this paradigm, contrary to the findings after the acute

injection, no temporal correlation between changes in PeVN mRNA levels and peptide hypothalamic contents were observed. These data suggest that long-term BDNF administration strongly activate SRIH translation process which is probably associated to changes in the messenger turnover. Long continuous infusion of BDNF could also lead to modifications in the chemical environment and thereby induce structural rearrangements of the target brain regions investigated. Indeed, the balance between excitatory and inhibitory neurotransmission (Tapia-Arancibia *et al.* 2004) can be modified by a long-lasting exposure to BDNF. For instance, some neuropeptides can be hypersecreted as a consequence of the food intake decrease. That could be the case of vasoactive intestinal peptide (VIP) since on the one hand, it is involved in the food intake regulation (Tachibana *et al.* 2003) and on the other hand, it could be stimulated by BDNF (Villuendas *et al.* 2001) resulting in increased SRIH secretion (Tapia-Arancibia & Reichlin 1985, De Los Frailes *et al.* 1991, Villuendas *et al.* 2001).

The BDNF induced-body weight loss observed here has been notably explained by a decrease in food intake (Pelleymounter *et al.* 1995, Siuciak *et al.* 1996). Among several other neuropeptides proposed (Wisse *et al.* 2003), hypothalamic SRIH (Feifel & Vaccarino 1994, Scacchi *et al.* 2003) and VIP (Tachibana *et al.* 2003) could also be involved in the weight loss given their roles in food intake regulation. In anorexic humans, a low somatostatinergic tone has been related to the weight loss (Stoving *et al.* 2002).

Besides, BDNF i.c.v. administration induces an increase in corticotropin releasing hormone (CRH) content and CRH mRNA level in the PVN (Givalois *et al.* 2004). CRH is known as a potent anorexigenic agent (Kalra *et al.* 1999). Since interactions between SRIH and CRH have been described (Aguila & McCann 1985, Spada *et al.* 1990, Hisano & Daikoku 1991), the BDNF/SRIH interaction could be exerted via CRH-mediation. Finally, we have also observed that long-term BDNF administration increases daily locomotor activity and temperature rhythms (Naert *et al.* unpublished observation). All of these actions may contribute to explain the weight loss measured in chronic BDNF-treated rats.

BDNF activation of TrkB receptors involves multiple second-messenger pathways that may underlie the described effects. It has been reported that cAMP responsive element binding (CREB) protein is a major mediator of neuronal neurotrophin responses (Finkbeiner *et al.* 1997). On the other hand, we previously showed that cAMP increased SRIH biosynthesis in experiments conducted with rat hypothalamic neurons or in NIH-3T3 fibroblast cells transfected with the rat SRIH gene (Montminy *et al.* 1986). Within the SRIH gene, a cAMP-responsive element has also been detected (Montminy *et al.* 1986). CREB is probably one signaling pathway by which BDNF activates SRIH gene expression in hypothalamic neurons.

In summary, our results show for the first time an *in vivo* regulation of SRIH hypothalamic neurons by BDNF in adult normal rats, which extend and confirm our previous *in vitro* data. Apart from a regulatory role in the food intake process above discussed, another physiological role of the BDNF/SRIH interaction might be found in the stress regulation. Indeed, as SRIH is massively released in the portal blood following application of different stressors, BDNF could contribute to the reconstitution of intracellular SRIH stocks once they have been exhausted by the strong stress demand.

Acknowledgements

This work was supported by the CNRS (France) and by grants from NARSAD (Independent Investigator Award 2002, USA) to LG, from Fondation Cino del Duca to S A

Figure 5 Effects of continuous BDNF infusion (12 µg/day per rat for 14 days) on hypothalamic SRIH. (A) illustrates the mean optical density (OD) and (B) the mean positive cell number/section of the SRIH mRNA signal in the PeVN of control and BDNF-treated rats. The results are expressed as means $\pm$ S.E.M. of the percent of control (considered as 100%), with $n=4-5$ rats per experimental group. ** $P<0.01$ versus control group. (C) shows hypothalamic SRIH contents in control and BDNF-treated rats. The results are expressed as means $\pm$ S.E.M. with $n=5-8$ rats per experimental group. * $P<0.05$ versus control group.

and from MENRT (France) to G N. The authors thank Regeneron Pharmaceuticals Inc. (USA) for the supply of BDNF and Mr E Savary for technical assistance. The authors declare that there is no conflict of interest that would prejudice the impartiality of this scientific work.

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Received 8 December 2005

Accepted 12 December 2005