Decreased serotonin transporter immunoreactivity in the human hypothalamic infundibular nucleus of overweight subjects

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Decreased serotonin transporter immunoreactivity in the human hypothalamic infundibular nucleus of overweight subjects

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INTRODUCTION

The brain serotonin (5-HT) system is known to be involved in the regulation of food intake and body weight. Several animal experimental studies manipulating endogenous serotonin synthesis, availability and metabolism, have made it clear that there is a negative relationship between the level of brain serotonin and food intake, in which increasing serotonin (by inhibiting reuptake or activating serotonin receptors post-synaptically) inhibits food intake (Lam et al., 2010). In addition, over the last few years evidence has accumulated on the central serotonin system’s involvement in peripheral glucose metabolism (Lam and Heisler, 2007). A more recent study reports that serotonin transporter (SERT) knockout mice are obese and present pre-diabetic symptoms such as glucose intolerance and insulin resistance (Chen et al., 2012). In peripheral glucose metabolism (Lam and Heisler, 2007).

Several neuropeptides related to both feeding behavior and glucose metabolism have been localized in the ARC/IFN. Like has been shown for rodents, Neuropeptide Y (NPY), agouti-related protein (AGRP) and α-melanocyte stimulating hormone (αMSH) are localized in the human IFN (Goldstone et al., 2002; Alkemade et al., 2012; Saderi et al., 2012). When these neuropeptides are injected into the rodent brain, NPY and AGRP increase feeding, reduce insulin sensitivity, and increase glucose production. On the other hand, αMSH decreases food intake and increases insulin action, thus reducing glucose production and enhancing glucose uptake (for review: Arble and Sandoval, 2013). More recently we investigated the expression of these neuropeptides in the human IFN in relation to body mass index (BMI), and type 2 diabetes
We found that AGRP expression showed a U-shaped correlation with BMI, and that NPY expression was lower in overweight and obese subjects, whereas αMSH revealed no relation to BMI.

Although it is clear that serotonin has an effect on feeding behavior and glucose metabolism in the rodent hypothalamus, data on human hypothalamus are scarce. What has been shown in humans is that polymorphisms related to serotonergic signaling are associated with BMI (Heisler and Tecott, 1999; Sookoian et al., 2008), however this does not provide evidence for hypothalamic serotonin as serotonin signals throughout the body and also widely throughout the brain. We did recently show that hypercaloric high fat high sugar snacking reduces diencephalic SERT (Koopman et al., 2013). Although diencephalon contains hypothalamic, it remains to be determined whether SERT containing cells/fibers can be identified in the human IFN and other hypothalamic nuclei. If identified, the question arises which individual nuclei contain the densest SERT staining and how SERT staining in the IFN relates to neuropeptides known to be involved in energy metabolism.

In this paper we determined the distribution of SERT immunoreactivity in the human hypothalamus using a post-mortem approach. In addition, we compared SERT staining in the IFN of subjects with a BMI < 25 to that of subjects with a BMI ≥ 25 kg/m². Finally, we describe immunofluorescent double-labeling of SERT with the IFN neuropeptides NPY, AgRP, and αMSH to identify cell types showing basket-like staining of SERT-immunoreactive fibers, suggestive of synaptic innervation.

MATERIALS AND METHODS

SUBJECTS

For Experiment 1 we investigated the distribution of SERT using systematic sampling over the entire rostro-caudal axis of the hypothalamus of 6 subjects (3 male) without neurological or psychiatric disease ranging in age between 67 and 86 years. Clinicopathological and relevant medication data are presented in Table 1.

For Experiment 2 in which we related SERT staining in the IFN to BMI, we studied post-mortem hypothalamic tissue of 11 overweight and obese (6 male, median BMI 30.5 (range: 25.0–39.5) kg/m²; median age 76 (range: 65–100) years) with 12 non-obese (5 male, median BMI 20.1 (range:15.2–24.2) kg/m²; median age 64 (range: 50–92) years) subjects. Clinicopathological and relevant medication data have been published earlier (Alkemade et al., 2012), and are presented in the Table 2.

In experiment 3 we aimed to identify the immunocytochemical nature of the cells showing basket-like SERT immunoreactivity, using brain material of the subjects described in Experiment 1. All brain material was obtained from The Netherlands Brain Bank at The Netherlands Institute for Neuroscience (director Dr. I. Huitinga) in accordance with the formal permissions for brain autopsy and for the use of human brain material and clinical information for research purposes.

HISTOLOGY

Brains were dissected at autopsy and the hypothalamus was fixed in 10% phosphate-buffered formalin at room temperature (RT) for 1–2 months. After dehydration in graded ethanol series, tissues were cleared in toluene, and embedded in paraffin. Coronal serial sections (6 μm) were cut over the entire rostro-caudal axis of the hypothalamus. For anatomical orientation, every 100th section was collected and mounted on chrome alum-gelatine coated glass slides and subsequently dried for 2 days at 37°C, followed by Nissl staining.

ANTIBODY CHARACTERIZATION

Mouse monoclonal anti-human SERT antibody was purchased from Millipore, MAB Technologies Inc. (Stone Mountain, GA; catalog no Mab5618). Antibody specificity has been reported before and was supported using Western blotting (Borman et al., 2000; Ramsey and Defelice, 2002; Serafeim et al., 2002; Henry et al., 2003). Rabbit polyclonal anti-human AGRP antibody was obtained from Phoenix Pharmaceuticals (Belmont, CA; catalog no. H-003-53). AGRP staining disappeared after pre-absorption with AGRP and was not affected by cross adsorption using the NPY peptide (Goldstone et al., 2002). The αMSH antibody was raised against the αMSH C-terminal, which is modified in αMSH free acid, and absent in ACTH, minimizing cross reaction with other POMC products. Staining was abolished after pre-absorption with the αMSH peptide (Elias et al., 1998).

IMMUNOHISTOCHEMISTRY

Histological and immunocytochemical processing was performed as described previously, with some minor modifications (Alkemade et al., 2012). For SERT-immunohistochemistry, a series of coronal sections at 100-section intervals over the entire rostro-caudal axis of the hypothalamus was mounted on Superfrost plus slides (Menzel Glaser, Germany) and dried for at least 2 days at 37°C. This resulted in 6–12 sections per subject. Next, antigen retrieval was performed using microwave treatment (Shi et al., 1997) and sections were stained using the avidine biotinylated complex method (Hsu et al., 1981), according to the following protocol: Sections were deparafinised in xylene and rehydrated through graded ethanol series. After rinsing in distilled water, the sections were washed in TBS (pH 7.6) and antigen retrieval was performed using microwave treatment (10 min 700 W) in TBS (pH 7.6). After adjustment to RT, sections were incubated in the primary antibody diluted 1:5000 in SUMI [supermix, 0.05 M Tris, 0.15 M NaCl, 0.5% Triton X-100 (Sigma, Zwijndrecht, The Netherlands), and 0.25% gelatine (Merck Darmstadt, Germany) (pH 7.6)] overnight at 4°C. The slides were rinsed in TBS (pH 7.6, 3 × 5 min) and incubated for 1 h at RT in the second antibody (biotinylated horse anti-mouse 1:400 in SUMI; Vector Laboratories, Burlingame, CA). After rinsing in TBS (pH 7.6, 3 × 5 min), the sections were incubated 1 h at RT in avidine biotinylated complex (1:800 in SUMI; Vector Laboratories, Burlingame, CA) and subsequently rinsed in TBS (pH 7.6, 3 × 5 min). Finally, sections were incubated in 0.5 mg/ml 3,3′-diaminobenzidine (Sigma) in TBS containing 0.2% ammonium nickel sulphate (BDH; Brunschwig, Amsterdam, The Netherlands) and 0.01% H₂O₂ (Merck) for approximately 15 min. The reaction was stopped in distilled water. The sections were dehydrated in ascending series of ethanol, cleared
Table 1 | Brain material experiment 1.

<table>
<thead>
<tr>
<th>Subject</th>
<th>Sex</th>
<th>Age</th>
<th>PMD</th>
<th>Fix</th>
<th>Cause of death, clinical diagnoses</th>
</tr>
</thead>
<tbody>
<tr>
<td>99101</td>
<td>M</td>
<td>69</td>
<td>19</td>
<td>41</td>
<td>Respiratory insufficiency, infarction distal brainstem, alcohol abuse, hypertension</td>
</tr>
<tr>
<td>00007</td>
<td>M</td>
<td>85</td>
<td>15</td>
<td>35</td>
<td>Myocardial infarction, bladder carcinoma</td>
</tr>
<tr>
<td>03054</td>
<td>M</td>
<td>67</td>
<td>4.5</td>
<td>50</td>
<td>Cardiogenic shock, multi organ failure, type 2 diabetes (insulin, metformin), COPD</td>
</tr>
<tr>
<td>98056</td>
<td>F</td>
<td>83</td>
<td>5</td>
<td>41</td>
<td>Respiratory insufficiency, colon carcinoma, type 2 diabetes (no data available on medication), cataract, arthrosis</td>
</tr>
<tr>
<td>95016</td>
<td>F</td>
<td>86</td>
<td>13.5</td>
<td>30</td>
<td>Decompensatio cordis, type 2 diabetes (tolbutamide), angina pectoris, nephropathy, retinopathy</td>
</tr>
<tr>
<td>01005</td>
<td>F</td>
<td>76</td>
<td>20</td>
<td>36</td>
<td>Respiratory insufficiency, Non-Hodgkin lymphoma, hypertension, basal cell carcinoma</td>
</tr>
</tbody>
</table>

Fix, duration of fixation in days; PMD, post-mortem delay in hours.

Table 2 | Brain material experiment 2.

<table>
<thead>
<tr>
<th>Subject</th>
<th>Sex</th>
<th>Age</th>
<th>PMD</th>
<th>Fix</th>
<th>BMI</th>
<th>Cause of death, clinical diagnoses</th>
</tr>
</thead>
<tbody>
<tr>
<td>09102</td>
<td>F</td>
<td>92</td>
<td>7</td>
<td>51</td>
<td>15</td>
<td>Cachexia, rectal carcinoma, old CVA</td>
</tr>
<tr>
<td>98072</td>
<td>M</td>
<td>79</td>
<td>17</td>
<td>31</td>
<td>17.0</td>
<td>Pneumonia, COPD, CVA, renal insufficiency</td>
</tr>
<tr>
<td>98091</td>
<td>M</td>
<td>50</td>
<td>8</td>
<td>59</td>
<td>18.9</td>
<td>Bronchopneumonia, sepsis, hepatocellular carcinoma, pneumococcus infection</td>
</tr>
<tr>
<td>09075</td>
<td>F</td>
<td>88</td>
<td>7</td>
<td>44</td>
<td>19.6</td>
<td>Respiratory insufficiency, COPD, cardiac insufficiency, PTCA, adenocarcinoma</td>
</tr>
<tr>
<td>91205</td>
<td>F</td>
<td>65</td>
<td>9:30</td>
<td>10</td>
<td>20.0</td>
<td>Cardiac failure, mammary carcinoma</td>
</tr>
<tr>
<td>10008</td>
<td>M</td>
<td>81</td>
<td>4:50</td>
<td>53</td>
<td>20.3</td>
<td>Angina pectoris, hypercholesterolemia, adenocarcinoma, arthritis</td>
</tr>
<tr>
<td>08052</td>
<td>F</td>
<td>62</td>
<td>8</td>
<td>73</td>
<td>21.0</td>
<td>Respiratory insufficiency, renal cell carcinoma, metastasized</td>
</tr>
<tr>
<td>98161</td>
<td>M</td>
<td>63</td>
<td>2:30</td>
<td>108</td>
<td>23.8</td>
<td>Respiratory insufficiency, squamous cell carcinoma of the oropharynx, metastasized</td>
</tr>
<tr>
<td>98030</td>
<td>M</td>
<td>88</td>
<td>7</td>
<td>44</td>
<td>19.6</td>
<td>Respiratory insufficiency, COPD, cardiac insufficiency, PTCA, adenocarcinoma</td>
</tr>
<tr>
<td>10008</td>
<td>F</td>
<td>60</td>
<td>7</td>
<td>53</td>
<td>19.2</td>
<td>Cardiac failure, angina pectoris, old CVA</td>
</tr>
<tr>
<td>97060</td>
<td>M</td>
<td>65</td>
<td>59:30</td>
<td>36</td>
<td>20.7</td>
<td>Myocardial infarction, unsuccessful resuscitation, atrial fibrillation, heart failure</td>
</tr>
<tr>
<td>00072</td>
<td>M</td>
<td>78</td>
<td>18</td>
<td>45</td>
<td>39.5</td>
<td>Renal insufficiency, atrial fibrillation, heart failure, dehydration</td>
</tr>
</tbody>
</table>

Fix, duration of fixation in days; PMD, post-mortem delay in hours; BMI, body mass index in kg/m².

Double-labeling by immunohistochemistry

To identify cell types showing basket-like staining of SERT-immunoreactive fibers in the IFN, we performed immunofluorescent double staining of respectively SERT with αMSH (1:20000), NPY (1:1000), and AGRP (1:3000) in hypothalamic sections containing the IFN of all 6 subjects.

After overnight primary antibody incubation at 4°C the slides were rinsed in TBS (pH 7.6, 3 × 5 min) and incubated in the secondary antibodies (biotinylated horse anti-mouse 1:400 in SUMI; Vector laboratories, Burlingame, CA) for 1 h at RT. Following rinsing in TBS (pH 7.6, 3 × 5 min), the sections were incubated 1 h at RT in avidine biotinylated complex (1:800 in SUMI; Vector Laboratories, Burlingame, CA), subsequently rinsed in TBS (pH 7.6, 3 × 5 min) and incubated in biotinylated tyramide (1:750 in SUMI, 0.01% H2O2 (Merck, Darmstadt, Germany)) for 15 min at RT followed by rinsing in TBS (pH 7.6, 3 × 5 min). SERT was detected in green by streptavidin-Alexa488 (1:1000; Invitrogen, Eugene, Or). The other peptides were visualized in red by respectively anti-rabbit Alexa594 (1:1000; Invitrogen, Eugene, Or) for NPY or AGRP and anti-sheep Alexa594 (1:1000; Invitrogen, Eugene, Or) for αMSH. This fluorochrome-conjugated antibody incubation was performed for 1 h at RT, followed by overnight incubation at 4°C.
Vectorshield with DAPI (Vector laboratories, Inc, Burlingame, CA) was used for nuclear staining and cover slipping. The sections were stored under dark conditions at 4°C until further analysis. Colocalization was assessed by visual inspection.

**QUANTITATIVE ANALYSIS**

SERT immunoreactivity was quantified using an unbiased masking procedure of the 3,3′-diaminobenzidine-Ni precipitate (Alkemade et al., 2012). For quantification of the immunoreactive signal, gray values of the DAB-Ni precipitate in the IFN were analyzed by computer-assisted densitometry using Image pro (Media Cybernetics, Silver Spring) and software developed at the Netherlands Institute for Neuroscience. Every 100th section containing the IFN was analyzed and estimates of the immunoreactivity were made by averaging the signal density of the 3 sections that showed the highest signal, resulting in arbitrary units (a.u.) (Alkemade et al., 2003).

**STATISTICAL ANALYSIS**

The data of the second experiment were analyzed with SPSS for Windows, version 19.0 (SPSS Inc. Chicago, Illinois, USA). Group differences in numerical variables were evaluated using the Mann-Whitney U-test for not normally distributed parameters. A stepwise linear regression analysis was performed to investigate the effects of possible confounding factors. The statistical significance level for all analyses was set at \( p < 0.05 \) (two-sided).

**RESULTS**

Many SERT-immunoreactive fibers and few scattered SERT-positive basket cells were found throughout the entire hypothalamus, except in the white matter tracts (Figure 1A). Basket-like staining was defined as aggregation of staining surrounding neuronal cell bodies, creating a basket-like appearance, suggestive of synaptic innervation. The general distribution of SERT was comparable in all subjects studied (Figure 1A). A strong inter-individual variation in staining intensity was observed. SERT positive fibers were present throughout the entire hypothalamus, with a denser network of SERT-immunoreactive fibers in the perifornical area and in close proximity to the anterior commissure. A plexus along the ependyma of the third ventricle wall also showed strong SERT-immunoreactivity (Figure 1B). In addition, the highest fiber density was observed in the IFN, and the suprachiasmatic nucleus, which is the central biological clock of the human hypothalamus (Figures 1C,D). In these areas, cell bodies and capillaries were directly surrounded by clusters of SERT-immunoreactive fibers, highly suggestive of SERT-positive nerve endings in contact with SERT-negative perikarya and capillaries (high power inserts of Figures 1C,D). The supraoptic nucleus, paraventricular nucleus, lateral tuberal nucleus, and tuberomamillary nucleus contained relatively small numbers of SERT-immunoreactive fibers. No differences were observed between males and females.

SERT-staining was observed in the IFN of all subjects. IFN SERT-immunoreactivity was lower in overweight subjects than in non-overweight subjects (\( p = 0.036 \)) (Figure 2A). An example of the difference between overweight and non-overweight subjects is illustrated in Figures 2B,C. A stepwise linear regression analysis performed for the factors age, sex, fixation duration, and post-mortem delay did not reveal any significant results.

To further characterize the cell types showing basket-like staining of SERT-immunoreactive fibers in the IFN, we performed immunofluorescent double staining of SERT with NPY, AGRP, or αMSH on sections containing the IFN. SERT-immunoreactive fibers were present at all levels in all studied subjects. Many single-labeled cells expressing NPY, AGRP and αMSH-immunoreactive cells were found in the IFN in all subjects. A very small minority of neurons immunoreactive for αMSH or AGRP showed basket-like SERT staining. As SERT showed no clear colocalization with NPY, neurons showing basket-like SERT staining remained largely unidentified (Figures 3, 4). Of note, the labeling of SERT in the IFN was pre-dominantly localized within the IFN, somewhat lateral to the NPY, AGRP-, and αMSH-immunoreactive cells. A number of these cells appeared to be strongly activated, as indicated by the presence of two nucleoli. Together, our results indicate that SERT axons project to a minority of αMSH and...
**DISCUSSION**

We showed that SERT protein is extensively expressed in the human hypothalamus. Moreover, one of the nuclei most heavily innervated was the SCN, which is in agreement with the distribution of SERT immunoreactivity reported in rodents (Legutko and Gannon, 2001) as well as in human and non-human primates (Moore and Speh, 2004; Emiliano et al., 2007). Interestingly, when analysing SERT protein in post-mortem sections of the IFN of overweight and obese subjects and comparing them to lean individuals, the amount of SERT protein was clearly reduced in obesity. This is in line with a large number of knockout and transgenic mouse studies, which, collectively, show an inverse relationship between brain serotonin signaling and food intake, as well as with pharmacological studies targeting serotonin signaling in rodents (for review: Lam et al., 2010). We cannot exclude that BMI in these subjects was influenced by severe illness, including malignancies such as adenocarcinomas. In addition, some subjects received antidiabetic medication. Immunoreactivity for serotonin has been reported before in the rodent ARC, where stained fibers were found in the immediate vicinity of capillaries and neurons (Warembourg and Poulain, 1985). Surprisingly, our double-labeling study indicates that these SERT immunoreactive fibers in the IFN only partly belonged to AGRP or αMSH basket cells. Interestingly, the majority of SERT fibers was located lateral to currently unidentified subgroups of neurons in the IFN.

AGRP neurons, as well as to currently unidentified subgroups of neurons in the IFN.
to the neuronal populations expressing peptides NPY/AGRP and αMSH which could point to a role for serotonin in a different (unidentified) neuronal population. Yet from rodent studies there are indications of a close relationship between serotonin and NPY in eating behavior. NPY/AGRP neurons receive serotonin input (Guy et al., 1988; Heisler and Tecott, 1999), and NPY mRNA is increased by administration of exogenous serotonin (Choi et al., 2006). Pharmacological inhibition of serotonin signaling reduced the feeding effect of NPY administration (Bendotti and Samanin, 1987; Grignaschi et al., 1995; Lam et al., 2010).

Interestingly, another area with dense innervation of SERT fibers was the suprachiasmatic nucleus (SCN), the area in the brain where the biological clock resides. This finding on dense innervation is in agreement with literature showing serotonin as regulator of the circadian phase. This innervation is in agreement with literature showing serotonin as regulator of the circadian phase. This innervation is in agreement with literature showing serotonin as regulator of the circadian phase.

ACKNOWLEDGMENTS
Brain material was obtained from the Netherlands Brain Bank (director I. Huitinga). We wish to acknowledge Bart Fisser for his excellent technical assistance. Anneke Alkemade received a VENI-grant (nr 916.86.020), and Susanne E. La Fleur was supported by a VIDI-grant (nr 917.96.331) of the Netherlands Organization for Health Research and Development. Anke J. Borgers and Karin E. Koopman were supported by PhD fellowship grants awarded by the AMC Executive Board.

REFERENCES


