A morphometric study of the preoptic area of the guinea pig

K. Bogus-Nowakowska, A. Robak, S. Szteyn, M. Równiak, B. Wasilewska, J. Najdzion

Department of Comparative Anatomy, University of Warmia and Mazury in Olsztyn, Poland

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The aim of the study was to provide the topography and morphometric characteristics of the preoptic area (POA) of the guinea pig. The study was carried out on the brains of sexually mature guinea pigs of both sexes. A uniform procedure was followed in the study of the paraffin-embedded brain tissue blocks of males and females. The blocks were cut in the coronal plane into 50 μm sections and stained according to the Nissl method. The guinea pig POA consists of four parts: the medial preoptic area (MPA), lateral preoptic area (LPA), periventricular preoptic nucleus (PPN), and median preoptic nucleus (MPN). The topography and general structure of POA parts are similar in males and females. However, the PPNa cells of females are more intensely stained and are more densely packed than the PPNa cells of males. For morphometric analysis, the MPA and LPA as well as PPN and MPN were considered respectively as uniform structures, namely MPA-LPA and PPN-MPN. The statistical analysis showed that the volume of the PPN-MPN was larger in males than in females, whereas the MPA-LPA volume did not differ between the sexes. Moreover, the numerical density and the total number of neurons were statistically larger in males than in females in both the MPA-LPA and PPN-MPN. The parameters describing POA neurons were larger for MPA-LPA neurons in comparison with the PPN-MPN neurons. However, in this respect no sex differences were observed in both studied complexes. (Folia Morphol 2010; 69, 1: 15–23)

Key words: morphometry, male, female, sex differences

INTRODUCTION

The preoptic area (POA), which is sometimes described as a part of the telencephalon, seems to be structurally and functionally continuous with the hypothalamus. It sends and receives input to and from many hypothalamic nuclei [2, 21, 28]. The preoptic area influences behavioural, neuroendocrine, and autonomic nervous system activity associated with reproduction. It controls male sexual behaviour [23, 24, 26] and plays a critical role in regulating the cyclic release of pituitary gonadotropins in females [36]. Furthermore, it takes part in the regulation of parental behaviour [33].

Since Raisman and Field [29] reported sex differences related to the synaptic organization of the POA in the rat, and Gorski et al. [17] described the sexually dimorphic nucleus of the rat preoptic area, there have been numerous reports dealing with the POA in mammals. Although a large amount of data concerning the POA exists, most of them refer to the endocrine, behavioural, and biochemical aspects. To our knowledge, a significant deficiency exists in the field of quantitative data on the POA in the available literature. There is no data, for example, dealing with the volume, total cell number of the whole POA, or even the particular POA nuclei. Some mor-
phometric studies, usually concerning different aspects of sexual dimorphism, were conducted only in chosen nuclei, subnuclei, or even in smaller parts. For instance, volumetric analyses of components of the medial preoptic region were done in the rhesus monkey [8] and in the ferret [12]. The volume and number of cells were measured only in the dorsocentral portion of the medial preoptic anterior hypothalamus in the macaque [38] and in the sexually dimorphic nucleus in the human brain [37]. The interstitial nuclei of the preoptic-anterior hypothalamic area were quantitatively analyzed in humans [1, 11]. Some more data refer to the morphometric parameters of the POA in rodents. For example, quantitative analysis of the anteriorly and centrally placed subnuclei of the medial preoptic nucleus [9] and the sexually dimorphic region of the medial preoptic area (MPA) [20] was performed in the guinea pig, and in the magnocellular subdivision of the medial preoptic nucleus in the hamster [19]. The total cell number [13] and the cell density per unit area [17] were estimated in the rat sexually dimorphic nucleus of the POA.

In the existing literature almost all the quantitative data, although not numerous, refer only to chosen components of the preoptic region, and there is a lack of studies concerning such parameters as, for example, volume, number of neurons, or their size in all POA components in the guinea pig. Moreover, some reports related to the dimorphic region of the POA, for example in rodents [4, 9, 20] and humans [25, 36], present divergent results. Therefore, taking into consideration all the above-mentioned reasons, the aim of our study was to provide a topography and the morphometric characteristics of the guinea pig preoptic region in males and females.

MATERIAL AND METHODS

The study was performed on the brains of sexually mature guinea pigs (Cavia porcellus). The material consisted of 10 brains, namely 5 of males and 5 of females. The study was conducted according to the rules of the Local Ethical Committee of the University of Warmia and Mazury. All efforts were taken to limit animal suffering and the number of animals used to get reliable data. The male and female brains were treated according to the same procedure. All animals were given a lethal dose of Nembutal (80 mg/kg) intraperitoneally and were then decapitated. The brains were removed from the skulls and placed into 4% neutralised formalin for immersion fixation for at least three months. Then the brains were embedded in paraffin and cut on a Leica microtome (Germany) into 50 μm sections in the coronal plane. The sections were stained with cresyl violet according to the Nissl method. The preparations were analyzed topographically and morphometrically. The morphometric analysis was carried out using a calibrated image analysis system which consisted of a computer with morphometric software (Multi-Scan 8.2, Computer Scanning Systems, Poland) and a light microscope combined with a digital camera (CM40P, VideoTronic, Germany). Brain sections were coded so that the analyses were done without knowledge of the specimen sex. Each series of (animal) brain sections was treated uniformly according to the following procedure.

Computer reconstructions of the preoptic area

Microscope images (512 × 512 pixels each) of each of the brain sections were digitally recorded using a camera combined with a microscope and a computer. The recorded images from one section were joined to form one digital section that contained the preoptic area and neighboring structures. All sections containing the POA were digitally reconstructed.

Volumetric analysis. The volume of the POA parts was calculated according to the formula described by deVito et al. [14]. The total volume (V0) of a structure consists of the sum of the volumes of all sections (Vn) containing the structure.

\[ V_0 = \sum V_n \]

The volumes (Vn) were calculated according to the following formula:

\[ V_n = \frac{Z_n}{3} \times \left( A_n + A_{n+1} + \sqrt{A_n \times A_{n+1}} \right) \]

\( A_n \) — area of the examined structure on the nth section; \( A_{n+1} \) — area of the examined structure on the \( n + 1 \) section; \( Z_n \) — distance between the two sections.

In the digital sections, the boundaries of the preoptic structures were outlined on the monitor with a cursor.

Because the lateral side of the MPA adjoins the medial lateral preoptic area (LPA) side and the cell structure of the two areas is similar, it is very difficult to outline this boundary in cresyl violet stained sections. For these reasons, the MPA and LPA were considered as one part — MPA-LPA. Similarly, the dorso-lateral border of the periventricular preoptic nucleus...
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The preoptic area of the guinea pig closely adjoins the median preoptic nucleus (MPN), and these nuclei cannot be distinguished from each other, so in the morphometric analysis they were also considered as one complex — PPN-MPN. The borders of the structures are shown in Figure 1. After outlining the MPA-LPA and the PPN-MPN boundaries their areas were calculated using a computer equipped with Multi-Scan 8.2 software.

Numerical density. The number of neurons in 1 mm$^3$ was estimated according to the optical disector method described by West and Gundersen [40].

Total number of neurons. The total number of neurons in the morphometrically examined POA parts was calculated according to the formula described by West and Gundersen [40]. The density and numbers of cells in the manuscript and in the graphs are presented in thousands.

\[ N_0 = V_0 \times N_{mm^3} \]

$V_0$ — volume of structure; $N_{mm^3}$ — numerical density of structure.

Morphometric parameters of neurons. The analysis of neuronal parameters was carried out with a 40× objective lens, and every second section was analysed for each individual: 5 females and 5 males. For each of the 10 individuals, more than 2000 neurons were measured. The first section within a series was selected randomly. Perikarya with distinct nuclei were outlined digitally and measured using a calibrated system and image-analyzing Multi-Scan 8.2 software. The following neuronal parameters were measured: area of soma, soma circumference, soma length (long axis), and soma width (short axis).

Statistical analysis

The data were analysed by using Statistica 6.0 software. The data are presented as mean value ± standard error of the mean. Significant differences between the sexes were determined using a one-way analysis of variance. The comparison of two means was made using Student’s t-test. The statistical significance level was set at $p < 0.05$.

RESULTS

The preoptic area of the guinea pig is situated in front of the hypothalamus, bilaterally from the supraoptic recess of the third ventricle, and reaches the anterior hypothalamus. It consists of four parts: the medial preoptic area, the lateral preoptic area, the median preoptic nucleus, and the periventricular preoptic nucleus (Figs. 1, 2). The most rostral part of the POA is made up of the MPA, the anterior pole of which appears in front of the decussation of the optic nerves. Just behind
it is the lateral preoptic area. The similar cell structure of the LPA and the adjoining part of MPA, as well as their loose arrangement, cause these areas to be indistinctly separated from each other. The areas constitute quite extensive cell groups that are arranged parallel to the anterior commissure, which, together with the nucleus of stria terminalis, create their dorsal border. Medially, the MPA is bound in front by the diagonal band of Broca, whereas in the middle and posterior sections it is bound by the PPN and MPN. From the lateral side, the MPA is restricted by the LPA, whereas the lateral border of the LPA is adjacent to the globus pallidus and substantia innominata. Ventrally, the areas are limited by the basal nucleus of Meynert. Small groups of cells — the posterior poles of the MPA and LPA — pass into the anterior and lateral hypothalamic area, respectively, without any distinct border on the level of the anterior sector of the supraoptic nucleus.

The PPN and MPN appear slightly behind the LPA anterior pole and are small clusters of tightly packed cells. The PPN is located ventrally to the MPA and laterally to the supraoptic recess, whereas the MPN is an unpaired band of grey matter placed in the middle of the POA. The PPN posteriorly enlarges in the lateral and dorsal direction taking a triangular shape. The base of the triangle lies parallel to the optic chiasm, the medial edge is adjacent to the third ventricle, and the lateral edge adjoins the MPA (ventral portion) and the MPN (dorsal portion). In the middle and posterior sections the PPN is divided into the medial (PPNa) and lateral parts (PPNb) (Figs. 1A, B). The PPNa cells are more intensely stained and are more densely packed than the cells of the PPNb. The PPN division is more distinct in the female than in the male.

The median preoptic nucleus forms a kind of PPN continuation in the dorsal direction. From the ventral side the MPN adjoins the PPN, from the lateral it is encircled by the MPA, and dorsally it reaches the fornix. Posteriorly, an elongated stripe of cells creating the MPN lengthens considerably in the dorsal direction where it connects the columns of the fornix. More caudally, the MPN is divided by the commissura anterior into two parts: dorsal and ventral. The dorsal part has the form of a triangle the base of which is located above and parallel to the commissura anterior, and its sides are bounded by the columns of the fornix. The ventral part has the form of a vertical strip, lying under the commissura anterior, which widens significantly ventrolaterally so that the cells of the MPN enclose the dorsal and lateral borders of the PPN. The similar cell structures of the MPN and PPN, as well as their adjacent location, cause the border between these nuclei to be indistinct. In the caudal sections the PPN and MPN decrease and are separated from each other. The MPN posterior pole forms the posterior pole of the POA and disappears at the anterior level of the periventricular nucleus of the hypothalamus.

**Morphometric analysis**

**Volume.** The mean overall volume of the whole preoptic area for males was $0.85 \pm 0.02 \text{ mm}^3$ compared to the mean POA volume in females of $0.79 \pm 0.02 \text{ mm}^3$. The results of volumetric analysis of the individual POA parts are shown in Figures 3 and 4. The analysis of variance (ANOVA) indicated that the mean volume occupied by the MPA-LPA (Fig. 3) was not statistically different between males and females ($p > 0.05$) whereas the PPN-MPN mean volume (Fig. 4) was significantly larger in males than in females ($p < 0.05$).

**Density of neurons.** The mean number of neurons in 1 mm$^3$ in the whole POA for males was $124 \pm 2.9$ compared to the mean number of $98 \pm 2.3$ in females. In both males and females the mean neu-
Neuronal density was higher in the PPN-MPN than in the MPA-LPA (Figs. 5, 6). Moreover, the numerical density was sex-dependent: neurons within the MPA-LPA as well as in the PPN-MPN (Fig. 5, 6) were significantly more densely packed in males than in females (p < 0.05).

Total number of neurons. The sex differences observed in the numerical density and in the volume correspond to the sex differences in the total number of POA neurons. The mean total neuron number of the whole POA in males was 87 ± 3.2 compared to 60 ± 1.6 in females. Thus, in the whole male POA there were approximately 31% more neurons than in the female POA. The statistical analysis indicated that the mean total neuron number of MPA-LPA (Fig. 7), as well as of the PPN-MPN (Fig. 8), was significantly larger in males than in females (p < 0.05).

Area of neuronal cell bodies. The morphometric parameters of neurons for the individual POA parts are shown in Table 1. In both sexes the mean neuronal area of the MPA-LPA was larger than the area of the PPN-MPN neurons. The statistical analysis indicated that the neuronal area of the two examined parts differed significantly (p < 0.05). However, there was no significant sex difference in the neuronal area in the MPA-LPA (p > 0.56) or in the PPN-MPN (p > 0.32).

Neuronal body circumference. The mean neuron circumference of the MPA-LPA was larger than the parameter of PPN-MPN neurons in both males and females (Table 1). The two parts of the POA differed significantly in terms of this parameter (p < 0.05). However, statistical analysis demonstrated no significant differences between males and females in the neuron circumference in the MPA-LPA (p > 0.60) or in the PPN-MPN (p > 0.35).

Neuronal body length. The mean neuron length of the MPA-LPA was always larger than the mean neuron length of the PPN-MPN in both sexes (Table 1).
The length of the MPA-LPA neurons differed in a statistically significant way from the length of PPN-MPN neurons (p < 0.05). The neuron length, as the remaining parameter describing POA neurons, did not differ significantly between the sexes either in the MPA-LPA (p > 0.09) or in the PPN-MPN (p > 0.15).

**DISCUSSION**

The general structure and topography of the guinea pig preoptic area is similar to that described in other rodents [4–6, 15], insectivores [31], and sheep [30]. Even though some authors present other POA divisions, it is possible to carry out a comparative analysis. For example, Broadwell and Bleier [6] distinguished the triangular nucleus of the mouse that probably corresponds to the dorsal part of the median preoptic nucleus of the guinea pig. On the other hand, the medial preoptic nucleus of the rat and mouse [5, 6] seems to correspond to part of the guinea pig MPA (present study). The PPN nucleus of the guinea pig in the middle and posterior sectors contains two subgroups: the medial PPNa and the lateral PPNb. The PPNa, which is more intensely stained and more densely packed in the female guinea pig, appears to correspond to the MPNa [9] and the medial preoptic nucleus of the guinea pig [4]. According to Bleier et al. [4], the medial preoptic nucleus of the female guinea pig, just as the PPNa in our study, is also more tightly packed in the female than in the same region of the male.

Our study is the first concerning the morphometry of the whole preoptic area in the guinea pig. Although a few morphometric reports of guinea pig POA providing some volumetric measurements are available, they concern only some nuclei, or even subnuclei, usually showing sexual dimorphism.

The volumetric analysis of the guinea pig POA revealed that the PPN-MPN complex differed significantly between the male and female, whereas the volume of the MPA-LPA did not show sexual differences in this parameter. In our study we did not dis-

**Figure 7.** Mean estimated total number of neurons ± SEM × 10³ within the medial and lateral preoptic area (MPA-LPA) in the male and female guinea pig; *p < 0.05, in comparison to the female group.

**Figure 8.** Mean estimated total number of neurons ± SEM × 10³ within the periventricular and median preoptic nuclei (PPN-MPN) in the male and female guinea pig; *p < 0.05, in comparison to the female group.

**Table 1.** Mean estimated parameters (± SEM) of neurons of the medial and lateral preoptic area (MPA-LPA) and the periventricular and median preoptic nuclei (PPN-MPN) in the male and female guinea pig

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<tr>
<td>MPA-LPA</td>
<td>Male</td>
<td>106.91 ± 2.41</td>
<td>41.00 ± 0.65</td>
<td>15.08 ± 0.31</td>
<td>9.97 ± 0.12</td>
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<tr>
<td></td>
<td>Female</td>
<td>110.00 ± 3.25</td>
<td>41.60 ± 0.72</td>
<td>15.00 ± 0.26</td>
<td>10.27 ± 0.16</td>
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<tr>
<td>PPN-MPN</td>
<td>Male</td>
<td>76.59 ± 1.83</td>
<td>34.25 ± 0.55</td>
<td>11.93 ± 0.21</td>
<td>8.75 ± 0.09</td>
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<tr>
<td></td>
<td>Female</td>
<td>80.02 ± 1.96</td>
<td>35.04 ± 0.5</td>
<td>12.17 ± 0.19</td>
<td>9.01 ± 0.10</td>
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tinguish a group of cells corresponding to the sexually dimorphic nucleus of the preoptic area (SDN-POA), which was described by Gorski and Harlan [18] in the rat as several times larger in males than in females. The cytoarchitectonic study of the sexually dimorphic nuclear complex of the medial preoptic-anterior hypothalamic area (SDNC-MPAH) in rodents (guinea pig, rat, hamster, mouse) showed that the entire configuration was similar in size and contours in the two sexes, but that the dimorphism of the region was related to patterns of cell distribution and density. The medial preoptic nucleus was larger and, except in the mouse, appeared to have higher cellular density in females than in males [4].

According to Hines et al. [20], the darkly staining portion of the medial preoptic area of the guinea pig is approximately four times larger in the male than in the female. However, Byne, and Bleier [9] observed sex differences in two components of the medial preoptic nucleus of the guinea pig. The anteriorly placed subnucleus corresponding to the SDN-POA of the rat [18] was twice as large in females as in males, and a centrally placed subnucleus had about ten-times greater volume in males than in females. On the other hand, Bleier et al. [4] identified a group of cells corresponding to the SDN-POA described by Gorski and Harlan [18] in the guinea pig only in the male medial preoptic nucleus while in the female such a cell group could not be distinguished. However, in the study of the guinea pig preoptic region [9] a cell subgroup that resembles the SDN-POA of the rat [18] was observed in 9 out of 20 females. In the study by Hines et al. [20], a group of cells corresponding to the rat SDN-POA was distinguished in the male and female guinea pig. It should be noted, however, that in the studies of Byne and Bleier [9] and Hines et al. [20] an anterior hypothalamic nucleus showing sexual differences has been described. In addition, such discrepancies between the observations may result from the differences in technical or histological procedures [4, 9, 20]. Apart from rodents, sexual dimorphism relating to the volume of the POA nuclei has also been reported in a number of other species, including the sheep [32], ferret [12], rhesus monkey [8], and humans [1, 25, 36].

The ovine sexually dimorphic nucleus (oSDN), which is a part of the medial preoptic area/anterior hypothalamus, is three times larger in volume in rams than in ewes [32]. In addition to the sex difference, the study of Roselli et al. [32] revealed that the oSDN is two-times larger in female-oriented rams than in male-oriented rams. Because the preoptic area is known to control the expression of the male sexual behavior such findings may suggest that naturally occurring variations in sexual partner preferences may be related to differences in brain anatomy [32].

Like in rodents, quantitative studies of the human preoptic region have also produced discrepant results. Swaab and Flier’s [36] analysis of the SDN revealed that the nucleus is larger in men than in women. Allen et al. [1] did not confirm these findings. Nevertheless, they found two other nuclei, the interstitial nuclei of the anterior hypothalamus INAH-2 and INAH-3, that were larger in males than in females. In INAH-1, which corresponds to the SDN of Swaab and Fliers [36], and INAH-4 Allen et al. [1] found no sexual dimorphism. The sex-related difference in the volume of the human INAH-3 was confirmed by LeVay [25] and Byrne et al. [10]. They stated that the human INAH-3 occupied a greater volume in males than in females. However, they did not find any sex differences in other parts of INAH. According to Allen et al. [1] and Byrne et al. [10], such discrepancies among the nuclei of the preoptic region may be attributed to methodological differences. For example, studies that detected no sexual dimorphism in INAH-1 [1, 25] were conducted with the use of much thicker sections than those of Swaab and Fliers [36] and Swaab and Hofman [37].

Larger volume of INAH-3 in men [10] is associated with an increase in the number of neurons within this nucleus, but not with sexual differences in neuron size or density. Our results support these findings to some extent. As regards the size of neurons, we agree with Byrne et al. [10], as neither in the MPA-LPA nor in the PPN-MPN were sex differences in any parameter describing the size of neurons observed, as in the POA neurons of the hamster [19]. However, Brown et al. [7] found that POA neurons located beneath the anterior commissure were significantly larger in females than in males of the 129SvEv strain mouse, but not in other examined strains. Furthermore, cells from another group of the POA were larger in males than in females of the C57BL/6J and SF-1 gene-disrupted wild-types.

Taking into consideration the number of neurons in the human INAH-3, we agree with Byrne et al. [10] that a higher number of neurons occurs in the male nucleus. Contrary to their findings, however, we found that the neuronal density differed between sexes, and there were more neurons in 1 mm² in the male than in the female guinea pig, both in the PPN-MPN and MPA-LPA. Thus, the increased number of
neurons within the studied complexes in male guinea pigs was related to both the increased volume and the increased neuronal density. Therefore, the neuron number was higher not only in the male PPN-MPN complex (the volume of which was larger in males than in females), but also in the male MPA-LPA of the guinea pig. Cherry et al. [12] found a significantly higher number of cells present per unit area of section in males than in females in the dorsal region of the ferret preoptic/anterior hypothalamic area on day E41, whereas on day E33 the density of cells in both sexes was similar. Our findings indicating that the total number of neurons was higher in the male preoptic area are similar to the results of studies on different nuclei within the preoptic region of the rat [18, 27], gerbil [22], hamster [19], and humans [36]. In hamster MPN [19], the same as in the MPA-LPA of the guinea pig, sex differences in the neuron number are compensated for by sex differences in cell density, although the volume is not sexually dimorphic.

According to Swaab and Hofman [37] and Swaab et al. [35], sex differences concerning the neuron number may vary in different stages of human life, and even in adulthood their magnitude does not remain constant. At the time of birth, the total cell number of the SDN-POA is similar in boys as it is in girls and increases equally in both sexes until the age of four years. From that age, the cell number starts to decrease in girls. In boys, the cell number of the SDN-POA remains stable until approximately 50 years of age, when it starts to decrease, which results in much less pronounced sex difference in the cell number in this period. In women, the second phase of marked cell loss takes place after the age of 70 [37]. Beck et al. [3] suggest that the structure of limbic brain regions is dynamic in adulthood and the process of neuronal aging and changes in sex hormone levels during aging seem to be instrumental in the changes occurring in dimorphic brain regions [34]. The latest study of the adult rat indicates considerable plasticity of the brain dimorphic regions in which circulating androgens are required to maintain soma size, but not regional volume in males. However, ovarian steroids maintain both soma size and regional volume in females [16].

The study carried out on the guinea pig confirms that the preoptic area is a dimorphic structure, although our attention was not focused on the dimorphic areas of the POA. The guinea pig, as a precocial animal [39], may provide a better model of particular developmental and neuroendocrine processes in primates than would other rodents. The study of sex differences in the brain may complement our understanding of the relationship between brain structure and function or behaviour and cognition. Sex differences in the brain are not only the basis for sex differences in reproduction, gender identity, and sexual orientation, but also sex differences in the occurrence of psychiatric and neurological diseases in adulthood and in age-related neurodegeneration [35].

REFERENCES


