Interactions between Kisspeptin Neurons and Hypothalamic Tuberoinfundibular Dopaminergic Neurons in Aged Female Rats

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Received August 22, 2016; accepted October 24, 2016; published online December 22, 2016

Kisspeptin neurons in the arcuate nucleus (ARC) regulate prolactin secretion, and are in physical contact with tuberoinfundibular dopaminergic (TIDA) neurons, which inhibit prolactin secretion. Prolactin levels in the blood are increased with advancing age in rats; therefore, we investigated the interactions with TIDA neurons and kisspeptin neurons in aged female rats (24 months of age), relative to those of young adult female rats (9–10 weeks of age). Plasma prolactin levels in the aged rats were significantly higher than those of young adult rats. Tyrosine hydroxylase (TH)-immunoreactive (ir) cell bodies and kisspeptin-ir nerve fibers were found in the dorsomedial ARC of both groups. The number of TH-ir cell bodies in the dorsomedial ARC did not differ significantly between groups. Additionally, no significant differences in the number of TH-ir cells in contact with kisspeptin-ir fibers was observed between groups. However, the number of kisspeptin-ir or Kiss1 mRNA-expressing cells in the ARC was significantly reduced in the aged rats compared with that of the young rats. These results suggest that the contacts between TIDA neurons and kisspeptin neurons are maintained after reproductive senescence, while production of kisspeptin in the ARC decreases significantly during aging.

Key words: kisspeptin, prolactin, tyrosine hydroxylase, aging, dopamine

I. Introduction

Prolactin plays an important role in reproduction, such as the promotion of lactation and reproductive behavior. In rodents, prolactin also has effects on luteal function [4]. The release of prolactin is stimulated by several factors such as thyrotropin-releasing hormone (TRH) and estrogen, and inhibited by dopaminergic neurons [5, 8]. The dopaminergic neurons that regulate prolactin secretion are located within the hypothalamic arcuate nucleus (ARC), and are subdivided into three groups: the tuberoinfundibular (TIDA), tuberohypophyseal, and periventricular hypophyseal dopaminergic neurons [4]. The cell bodies of TIDA neurons are found in the dorsomedial ARC and project to the median eminence. Therefore, dopamine secreted from TIDA neurons is released in the pituitary portal blood and inhibits prolactin release in lactotrophs through activation of D2-like receptors [4, 5]. The activity of TIDA neurons is regulated by several neurotransmitters, neuromodulators and peripheral hormones [4]. Inhibitory agents such as serotonin, noradrenaline and estrogen are able to promote prolactin release through a decrease in TIDA activity. However, stimulatory agents such as acetylcholine, vasopressin and prolactin are able to promote the activity of TIDA neurons, resulting in decreased prolactin secretion.

Kisspeptin is a neuropeptide with an important role in the regulation of ovulation and follicular development through stimulation of GnRH/luteinizing hormone (LH) release in mammals [18]. In rodents, kisspeptin neurons located in the ARC coexpress neurokinin B (NKB) and dynorphin; described as KNDy neurons, these cells are thought to be involved in the generation of pulsatile GnRH/LH secretions [12]. Furthermore, kisspeptin induces...
prolactin release through inhibition of TIDA neurons in the presence of estrogen [17], and its fibers are in physical contact with TIDA neurons [19]. Kisspeptin neurons also express prolactin receptors [11] and are suppressed by prolactin [2], suggesting that kisspeptin and prolactin levels are regulated in coordination with each other.

In rats, serum prolactin levels increase with age [13, 22]. Serum prolactin levels of aged male rats (24–26 months of age) are, therefore, higher than those of young male rats (2–3 months of age) [6]. Dopamine levels in pituitary stalk blood plasma samples from aged rats are lower than those of young rats; thus, the increased incidence of hyperprolactinemia observed with advancing age may be associated with a reduction in dopamine secretion from TIDA neurons. Expression of the kisspeptin gene in the ARC of middle-aged female rats is lower than that of young rats [9]. Based upon these findings, this study aims to investigate the interaction between kisspeptin and TIDA neurons in aged female rats using immunohistochemical and in situ hybridization analysis.

II. Materials and Methods

Animals

Young adult (8 weeks of age, Tokyo Laboratory Animals Science, Tokyo, Japan) and aged (24 months of age, supplied by the Tokyo Metropolitan institute of Gerontology, Tokyo, Japan) female Wistar rats were housed in a controlled (14-hr light/10-hr dark, 6:00 lights on) environment with free access to food and water. Young rats that had undergone at least two consecutive 4-day estrus cycles were deemed suitable for investigations. Young (diestrus) and aged rats were deeply anesthetized with sodium pentobarbital, and blood samples were collected from the cervical vein. After blood sampling, anesthetized rats were perfused through the heart with a physiological saline solution, followed by 4% paraformaldehyde (PFA) in a 0.1 M phosphate buffer (PB). The brain was removed from the skull and stored in 4% PFA solution overnight at 4°C. The following day, the brain was immersed in 0.05 M PB containing 30% sucrose at 4°C for 4 days to protect against damage during freezing. One aged rat was excluded from analyses owing to the presence of a visually abnormal pituitary. Rat brains were used for immunohistochemistry and in situ hybridization analysis. All experimental procedures involving live rats were performed according to NIH guidelines (Guide for the Care and Use of Laboratory Animals) and received ethical approval from the Nippon Medical School Committee on Animal Research.

Dual-labeling fluorescence immunohistochemistry with kisspeptin and tyrosine hydroxylase (TH)

Rat brains were sectioned along the coronal plane at a thickness of 40 μm using a cryostat (CM-3050-S, Leica, Wetzlar, Germany). Serial sections containing the ARC (from 1.72–4.36 mm posterior to the bregma [14]) were obtained. Every fourth section through the dorsomedial ARC of each animal was stained. These samples were washed in 0.1 M PB containing 0.9% NaCl and 0.3% Triton-X 100 (Wako, Osaka, Japan) (PBST), and incubated with blocking buffer (1% bovine serum albumin (Sigma, St Louis, MO), 5% normal donkey serum and 0.02% sodium azide in PBST) for 90 min, followed by incubation with anti-rat kisspeptin antibody (1:2000, mouse monoclonal antibody, provided by Dr. Ohtaki, Takeda Pharmaceutical, Osaka, Japan) [10] and anti-TH antibody (1:500, rabbit polyclonal antibody, Chemicon International Inc., Temecula, CA) diluted in 0.1 M PBST at 4°C for 2 days. After washing in 0.1 M PBS, the sections were incubated with Alexa Fluor 568-conjugated donkey anti-mouse IgG (1:500, Invitrogen, Eugene, OR) and Alexa Fluor 400-conjugated donkey anti-rabbit IgG (1:500, Invitrogen) diluted with 0.1 M PBS for 2 hr at room temperature. Sections were then washed in 0.1 M PBS, and mounted on glass slides using Vectashield (H-1400, Vector Laboratories, Burlingame, CA) as the mounting medium. Fluorescence images were obtained using confocal laser microscopy (Fluoview FV10i, Olympus, Tokyo, Japan). Two sections with similar distributions of TH-immunoreactive (ir) cells in the dorsomedial ARC were selected from the stained sections of each animal, and Z-stack images were obtained. Z-stack images of each brain slice were acquired at 1.2-μm intervals. Contacts between at least one TH-ir cell body and one kisspeptin-ir fiber were confirmed through visual inspection of images showing TH-ir cells. Numbers of TH-ir cells with, or without contact with kisspeptin fibers were counted in two sections from each animal. The total sum of the cell number was recorded (n=4–6 per group).

In situ hybridization

Kiss1 in situ hybridization for Kiss1 (the gene encoding kisspeptin) was performed as described previously [1]. Briefly, serial 50-μm coronal sections containing the ARC were obtained as described in the previous paragraph, and every fourth section through the ARC (from 1.72 mm to 4.36 mm posterior to the bregma) was analyzed. A Kiss1-specific digoxigenin (DIG)-labeled probe was used to conduct free-floating in situ hybridization. A DIG-labeled antisense RNA probe was synthesized from a full-length rat Kiss1 template cDNA (GeneBank accession #AY196983) [23] using a DIG RNA labeling kit (Roche Diagnostics, Mannheim, Germany). Every fourth section through the ARC was hybridized with the labeled probe. To visualize DIG labeling, sections were incubated with an alkaline phosphatase (AP)-conjugated anti-DIG1 antibody (1:1000, Roche Diagnostics) for 2 hr at 37°C and then placed in 4-nitroblue tetrazolium chloride/5-bromo-4-chloro-3-indolyl phosphate solution (Roche Diagnostics). Kiss1-expressing cells in the ARC were manually counted under a light microscope (BX51 microscope, Olympus, Tokyo, Japan) and the total cell number in brain sections from each animal was recorded (n=4 for each group).
Kisspeptin immunohistochemistry

Serial 40-μm thick sections were obtained for double staining. Every fourth section through the ARC (from 1.72 mm to 4.36 mm posterior to the bregma) was analyzed. These samples were incubated with anti-rat kisspeptin antibody (1:5000) for 24 hr at 4°C. Sections were washed and incubated with biotin-conjugated anti-mouse IgG (Histofine SAB-PO (M) kit, Nichirei Biosciences, Tokyo, Japan) for 2 hr, followed by incubation in avidin-biotinylated horseradish peroxidase (HRP) complex [Histofine SAB-PO (M) kit] for 2 hr. Immunoreactivity was visualized using 3′3-diaminobenzidine (DAB) mixed with 0.03% hydrogen peroxide. Kisspeptin-ir cells in the ARC were manually counted under a light microscope (BX51) and the total cell number in brain sections from each animal was recorded (n=4–6 per group).

Prolactin assay

Following blood sampling, plasma was isolated immediately using centrifugation and stored at −20°C until analysis using a prolactin assay. Plasma prolactin levels (n=5 for each group) were determined using a commercial kit (rat prolactin ELISA kit, cusabio biotech, Wuhan, China). Plasma samples from aged rats were diluted four-fold with distilled water owing to readings from undiluted samples exceeding the upper limits of the detection range.

Statistical analysis

All data are reported as mean±standard error of the mean (SEM). Statistical differences (P<0.05) between groups were determined using an unpaired Student’s t-test.

III. Results

Plasma prolactin levels in aged rats were significantly higher than those of young rats (Fig. 1). Kisspeptin-ir fibers and TH-ir cell bodies were investigated using immunofluorescence microscopy, and were observed in the dorsomedial ARC of both aged and young rats (Fig. 2A). The numbers of TH-ir cells did not differ significantly between groups (Fig. 2B). Furthermore, no significant differences in the number of TH-ir cells in contact with kisspeptin-ir fibers were observed (Fig. 2B). Kisspeptin-ir fibers were observed in the dorsomedial ARC of both aged and young rats, although the number of kisspeptin-ir cell bodies in the ARC was clearly lower in aged rats compared with young rats (Fig. 3). Moreover, significantly lower numbers of Kiss1 mRNA-expressing cells were observed using in situ hybridization in the ARC of aged rats compared with those of young adult rats (Fig. 3).

IV. Discussion

In rats, prolactin levels increase with age [6, 13, 22]. In this study, plasma prolactin levels in aged rats were also higher than those of young rats. TIDA neurons are known to regulate prolactin secretion [5]. Nonetheless, the number of TH-ir cells did not differ significantly between groups. This result is consistent with that of previous studies [15, 16]. No significant differences in both the number of TH-ir cells and also levels of TH mRNA expression in the hypothalamus were observed in aged, compared with young rats [15, 16]. However, the same study demonstrated that dopamine levels in the hypophysial portal blood of aged male rats are lower than those of young male rats [15]. Also, TH activity in the medial basal hypothalamus was reduced in aged female rats compared with young female rats [15]; therefore, the high levels of prolactin in the blood of aged rats may be induced by a decrease in TH activity rather than a decrease in the number of TH-expressing cells.

Kisspeptin-expressing neurons in the ARC are in contact with TIDA neurons [19, 20], and regulate prolactin secretion through inhibition of TIDA neurons [17]. In this study, kisspeptin-ir fibers were found in the dorsomedial ARC of both young and aged rats. In addition, the number of TH-ir cells in contact with kisspeptin-ir fibers did not differ between young and aged rats. Numbers of kisspeptin-ir or Kiss1 mRNA-expressing cells in the ARC were reduced in aged rats compared with young rats; therefore, kisspeptin appears to have accumulated in the terminals of fibers owing to a reduction in kisspeptin production in aged rats. It is difficult to clearly explain the mechanism of this result. It is a possible explanation that the age-related changes about the axonal transport, release mechanism and/or metabolic status of kisspeptin in the axon terminal may also be considerable. The further study on these possibilities is necessary.

Previous studies have shown that kisspeptin increases prolactin secretion through inhibition of TIDA neurons in the presence of estrogen [17]. In this study, kisspeptin expression decreased in aged rats with hyperprolactinemia. The reasons for this discrepancy are unclear. However, there is a need to take into account a short loop feedback system between kisspeptin and prolactin; kisspeptin neurons in the ARC express the prolactin receptor [11] and are...
inhibited by prolactin [2]. Therefore, kisspeptin may act as a trigger of hyperprolactinemia and is inhibited by the increase in serum prolactin levels in aging rats.

Some endogenous signaling molecules are able to stimulate prolactin secretion. TRH stimulates prolactin secretion from lactotrophs [8]. High levels of proTRH are expressed in the hypothalamic paraventricular nucleus (PVN). Levels of TRH mRNA expression in the hypothalamic PVN of male rats decrease with advancing age [3]; therefore, TRH levels may no longer be sufficient to stimulate prolactin secretion. Estrogen is also able to stimulate prolactin secretion, through inhibition of TIDA neurons, because TIDA neurons express the estrogen receptor α (ERα), and are inhibited by estrogen [5]. In aged female rats in an anestrus state, serum estradiol levels decrease compared with those of young rats [7, 21]. Therefore, estrogen may not be involved in prolactin secretion in aged rats.

In this study, the extent of kisspeptin production in the ARC of aged rats was reduced compared with that of young rats, although no significant differences in TH-immunoreactivity in TIDA neurons and kisspeptin-immunoreactivity in the fibers of the dorsomedial ARC were observed with aging. Therefore, contacts between kisspeptin and TIDA neurons seem to be maintained during aging. Kisspeptin neurons may affect not so much a number of TH cells as a release/metabolism of TH, which may be associated with regulation of prolactin release in aged animals. The present study of alterations in kisspeptin

Fig. 2. Immunohistochemical labelling of tuberoinfundibular dopaminergic (TIDA) neurons and kisspeptin fibers in the dorsomedial arcuate nucleus (ARC). (A) Double immunostaining for tyrosine hydroxylase (TH, green) and kisspeptin (red) in the dorsomedial ARC of representative rats. Upper panels show images from the ARC of young rats during the diestrus stage. Lower panels show images from aged rats. Bars=50 μm. (B) Number of TH-ir cells (left panel) and TH-ir cells in contact with kisspeptin-ir fibers (right panel) in the dorsomedial ARC. Values are expressed as mean±SEM. Numbers in each column represent the number of animals examined. The statistical significance of these differences was determined using an unpaired Student’s t-test.

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neurons with advancing age provides further information on the changes in reproductive function observed during aging, and during age-related hormonal alterations, such as the menopause.

V. Disclosure Statement

The authors have nothing to disclose.

VI. Acknowledgments

We are grateful to Takeda Pharmaceutical Company Ltd. for kindly providing rat Kiss1 cDNA. This work was supported by Grants-in-Aid for Scientific Research (No. 26460323 to H. O. and No. 15K18979 to K. I.) from the Japan Society for the Promotion of Science (JSPS) KAKENHI, and the Japanese Ministry of Education, Sports, Science.
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