Estradiol Upregulates Kisspeptin Expression in the Preoptic Area of both the Male and Female Rhesus Monkey (*Macaca mulatta*): Implications for the Hypothalamic Control of Ovulation in Highly Evolved Primates

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Key Words

Kisspeptin · Preoptic area · Luteinizing hormone surge · Sexual dimorphism

Abstract

The aim of this immunohistochemical study was to evaluate the distribution of kisspeptin neurons in the preoptic area (POA) of gonadally intact adult male and female rhesus monkeys, and to determine whether imposition of an estradiol (E₂)-positive feedback signal in the castrate male increased kisspeptin in the POA. Additionally, kisspeptin in the POA of the intact female was examined during an LH surge induced prematurely by E₂ administered in the early follicular phase. The number of kisspeptin neurons in the POA of males and females was similar. Immunoactive kisspeptin perikarya were not observed in the POA of castrate adult males, but such neurons in these animals were present within 12 h of imposing an increment in circulating E₂ concentrations that in a screening study conducted 4–6 weeks earlier had elicited an LH surge. As expected, premature induction of an LH surge by E₂ early in the follicular phase was associated with

upregulation of kisspeptin in the POA. These results represent the first description of immunoreactive kisspeptin cell bodies in the POA of the macaque brain and provide further support for the view that (1) kisspeptin neurons in the POA of the female monkey are a target for the positive feedback action of E₂ and (2) the hypothalamic mechanism which mediates this action of E₂ in primates is not subjected to perinatal programming by testicular testosterone. Moreover, our findings indicate that maintenance of the kisspeptin content in the POA of intact male monkeys requires the action of E₂, presumably generated by aromatization of testicular testosterone at the hypothalamic level.

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Introduction

Classic studies of rodents have demonstrated that, in these species, the preovulatory LH surge is generated by a neural signal that originates in the rostral hypothalamus, specifically in the preoptic area (POA), and is transmitted to the pituitary via the hypophyseal portal circula-

tion by a large discharge or surge of GnRH [1]. The initiation of this GnRH surge is triggered by the interaction of a circadian signal [2] originating from the suprachiasmatic nucleus and a positive feedback action of preovulatory levels of E_2 in the POA [1]. The mandatory role of the POA for the LH surge in rodents was elegantly demonstrated in the rat by Halász and Gorski [3], who showed that after surgical isolation of the POA from the mediobasal hypothalamus (MBH), females became acyclic and failed to ovulate. Similar studies of female rhesus monkeys were performed by Knobil's group [4]; however, in contrast to rodents, surgical isolation of the MBH in these animals did not consistently interrupt menstrual cyclicity and ovulation, suggesting that the POA is not essential for the positive feedback action of E_2 that generates the preovulatory LH surge in highly evolved primates.

Kisspeptin neurons in the anteroventral periventricular area and the periventricular nucleus of the POA are generally recognized as a major target for the positive feedback action of E_2 that underlies the GnRH surge in rodents [5, 6]. Interestingly, *KISS1* mRNA is also found in the POA of the female monkey [7, 8] and human [9], and elevated *KISS1* expression in this hypothalamic area has been described in the female rhesus monkey during the late follicular phase of the menstrual cycle at the time the preovulatory LH surge was to be anticipated [8]. Together, these findings suggest that the POA of the primate hypothalamus may play a role, albeit a redundant one, in mediating the positive feedback action of E_2 to elicit the preovulatory surge of LH.

In rodents, the POA mechanism that triggers the preovulatory LH surge in adult females is decommissioned in males by an action of testicular testosterone secreted during the early postnatal period of development [10]. As may have been predicted from the established role of POA kisspeptin neurons in triggering the LH surge during the estrous cycle, kisspeptin expression in the POA of the mouse and rat is sexually dimorphic, showing a dramatic female-dominant sex difference in the number of these peptidergic neurons [11, 12]. Not surprisingly, this sexual dimorphism is also determined by testicular testosterone secretion during early postnatal development, as demonstrated by the finding that in female rats a single injection of testosterone propionate on the day of birth leads to a characteristic male-like pattern of kisspeptin neurons in the adult POA [13].

In contrast to rodents, the preovulatory gonadotropin surge in primates is not 'masculinized' by testicular androgen exposure during perinatal development. Adult

male monkeys, castrated as adults, are able to generate LH surges in response to exogenous E_2 [14], and when such animals are exposed for 30–60 days to low follicular phase levels of E_2 before elevating levels of this steroid to produce a positive feedback signal, the induced LH surge is markedly amplified [15]. A sensational demonstration of the ability of the hypothalamic-pituitary unit of this male primate to respond to the positive feedback action of E_2 was provided by the finding that an ovarian fragment transplanted subcutaneously into a castrated adult male rhesus monkey exhibited repetitive ovulatory cycles with approximately normal periodicity [16]. Similarly, intact and gonadectomized adult men also respond to the positive feedback action of E_2 , as demonstrated by an LH surge elicited after 5–9 days of exposure to exogenous E_2 [17, 18].

Given the lack of sexual dimorphism in the neuroendocrine mechanism that generates the preovulatory LH surge in highly evolved primates, Watanabe et al. [19] conducted a study to determine if POA kisspeptin neurons are activated by E_2 in male as well as in female monkeys. To test this idea, castrate adult male Japanese macaques were implanted with E_2 -containing capsules to produce circulating levels of the steroid typical of those observed in the early follicular phase of the menstrual cycle. After 2 weeks of exposure to these relatively low levels of E_2 , the animals received an intramuscular injection of E_2 benzoate at a dose previously established to induce an LH surge in this model; *KISS1* mRNA levels in POA kisspeptin neurons together with c-Fos expression, a marker of neuronal activation, were upregulated in association with the estrogen-induced LH surge. While this observation suggests that E_2 activates POA kisspeptin neurons when an LH surge is elicited in male monkeys, it should be noted that the study design did not allow for the positive feedback action of E_2 (generated by the E_2 benzoate injection) to be separated from the effects of the 2 weeks of low-dose E_2 exposure.

The present study was conducted to extend the foregoing findings on the regulation by E_2 of kisspeptin neurons in the POA of the rhesus monkey. Specifically, we attempted to (1) compare the distribution of immunoreactive kisspeptin in the POA of intact adult male and follicular-phase female monkeys, (2) examine whether the premature induction of an LH surge by exogenous E_2 administration early in the follicular phase of the menstrual cycle evokes increased kisspeptin expression in the POA, and (3) determine whether imposition of a positive E_2 feedback signal in the castrate adult male monkey is associated with an increase in

kisspeptin immunoreactive elements in the POA. The use of immunohistochemistry – rather than in situ hybridization, as previously employed – to monitor the distribution and activity of *KISS1* neurons in the POA provides the first description of the location of the peptide encoded by the gene in this region of the monkey hypothalamus.

Materials and Methods

Animals and Hypothalamic Tissue

Hypothalamic tissue for this study was obtained from 3 different sources.

Source 1

Four adult female rhesus monkeys (*Macaca mulatta*) ranging from 5 to 8 years of age and weighing 5–9 kg, and 6 castrate adult males (8–11 years of age, 8–12 kg body weight) were used. The females had menstruated spontaneously for 2–4 months with cycles ranging from 29 to 40 days before the study. The males had been gonadectomized 3–7 months earlier for other purposes. All monkeys were maintained under a controlled photoperiod (lights on between 07:00 and 19:00 h) and temperature (approx. 21°C) in accordance with the National Institutes of Health Guidelines for the Care and Use of Laboratory Animals. At the completion of each experimental protocol (see Experimental Design), the animals were first sedated with ketamine hydrochloride (approx. 20 mg/kg body weight; Ketaject, Phoenix Scientific Inc., St. Joseph, Mo., USA) and deeply anesthetized with sodium pentobarbital (approx. 30 mg/kg body weight, i.v.; Nembutal sodium solution, Abbott Laboratories, North Chicago, Ill., USA). The thoracic cavity was exposed, the dorsal aorta clamped, and the head perfused transcardially at a flow rate of approximately 40 ml/min, initially with approximately 1 liter of physiological saline containing 2% sodium nitrite and 5,000 U of heparin/l, and immediately followed with 2–3 liters of 4% paraformaldehyde (PFA) in 0.1 M phosphate-buffered saline (PBS; pH 7.2). Following perfusion, the brain was removed from the cranium and the hypothalamus isolated. A parasagittal cut was made at approximately 4 mm on either side of the midline, and a horizontal cut was made immediately dorsal to the anterior commissure. The hypothalamic block was immersed in fixative for 1–2 h and was then transferred to a 30% sucrose solution (in 0.1 M PBS, pH 7.2) for at least 24 h at 4°C. Serial sections throughout the entire hypothalamic block were cut on a freezing microtome every 25 µm. The sections were placed sequentially into a train of 10 wells, with each well containing a sequence of sections collected at 250-µm intervals from the region of the organum vasculosum of the lamina terminalis (OVLT) to the mammillary bodies. The sections were stored in an antifreeze cryoprotectant solution at –20°C [20]. The foregoing experimental procedures were approved by the University of Pittsburgh Institutional Animal Care and Use Committee.

Source 2

Hypothalami from 3 hysterectomized rhesus monkeys in the follicular phase of the menstrual cycle ranging in age from 8 to 15 years (9–11 kg body weight) were obtained post mortem after the

animals had been venipunctured and euthanized with Beuthanasia-D (Henry Schein Animal Health, Dublin, Ohio, USA) and their brains immediately perfused with approximately 1 liter of saline containing 2% sodium nitrite and 5,000 U heparin/l followed by approximately 2 liters of 4% PFA in 0.1 M PBS (pH 7.2). Following removal of the brain from the cranium, the hypothalamus was postfixed in 4% PFA at 4°C overnight prior to immersion in 30% sucrose. Serial 25-µm sections throughout the entire hypothalamus were cut and collected as described above.

Source 3

Twenty-five-micrometer sections from 3 intact adult males (6–10 years old and 8–10 kg body weight) and 3 adult castrate males (4–14 kg body weight and gonadectomized at the age of 3–10 years) were obtained from our tissue bank. The animals had been previously used for studies of testicular function, and castration had been performed 1–5 months before perfusion and postfixation, which were conducted exactly as described for the animals obtained from the first source. The hypothalamic sections had also been cryopreserved in wells containing a sequence of sections collected at 250 µm from the OVLT to the mammillary bodies.

Experimental Design

Experiment 1: Patterns of Kisspeptin Immunoreactivity in POA Perikarya and the Distribution of These Neurons in Intact Males and Females

Hypothalamic tissue obtained from 3 hysterectomized adult females and 3 intact adult males was used. It was determined that the females were in the follicular phase of the menstrual cycle at the time of perfusion based on the following hormonal data obtained on the morning of perfusion: circulating LH levels ranged from 0.1 to 1.1 ng/ml, E₂ from 22.9 to 63.9 pg/ml, and progesterone from 0 to 0.3 ng/ml. Sequential series of sections collected every 250 µm through the POA from a male-female pair were processed together to minimize immunohistochemical variation.

Experiment 2: Immunoreactive Kisspeptin Expression in the POA Associated with an E₂-Induced LH Surge Prematurely Elicited in the Early Follicular Phase

Four intact females on day 3 of the menstrual cycle were implanted under sedation with ketamine hydrochloride with either blank Silastic capsules or E₂-containing Silastic capsules to produce circulating E₂ levels of 200–300 pg/ml (n = 2/group). The ability of this estrogen regimen to induce a premature LH surge in a control cycle was established for all 4 monkeys prior to the experimental cycle. The interval between control and experimental cycles ranged from 4 to 11 weeks. Blood samples were collected by femoral venipuncture under sedation with ketamine hydrochloride every 12 h for the duration of capsule implantation during both the control and the experimental cycles. Hypothalami were collected during the experimental cycle after 36 h of exposure to E₂. A sequential series of sections collected every 250 µm through the POA of each of the 4 monkeys were processed in a single immunohistochemical run.

Experiment 3: A Comparison of Immunoreactive Kisspeptin Neurons in the POA of Intact and Castrate Adult Male Monkeys and the Effect of E₂ Treatment on the Castrates

Six castrate males were implanted with E₂-containing Silastic capsules for 12 h (n = 3) or 48 h (n = 3) prior to perfusion, with the

Fig. 1. Photomicrographs ($\times 10$) of coronal hemisections of POA taken at the level of the AVPe from the same section of a male (**a**, **b**) and a female (**c**, **d**) monkey, and immunostained for kisspeptin using the primary antibody GQ2. **a**, **c** Without preabsorption of GQ2. **b**, **d** Preabsorbed with human kisspeptin-54. 3V = Third ventricle. Scale bars = 100 μ m.

aim of producing circulating E_2 levels of 200–400 pg/ml. Before the experiment, the ability of this E_2 treatment to induce an LH surge was established in all 6 monkeys. The time between the control and experimental E_2 challenges ranged from 4 to 6 weeks. Blood samples were collected by femoral venipuncture under sedation with ketamine hydrochloride every 12 h for the duration of capsule implantation during both the control and the experimental E_2 challenges. A sequential series of sections collected every 250 μ m through the POA from a castrate treated with E_2 for 12 and 48 h were processed together with a comparable series of sections from an intact and untreated castrate adult male in a single immunohistochemical run.

Antibodies and Blocking Peptides

The kisspeptin antibody used was a sheep anti-human kisspeptin-54 antibody (GQ2), kindly provided by Dr. Stephen Bloom (Imperial College London, Hammersmith Hospital, London, UK). The specificity of GQ2 has been previously validated for immunohistochemical studies of monkey hypothalami [21]. Biotinylated rabbit anti-sheep IgG (Vector Laboratories, Burlingame, Calif., USA) was used as the secondary antibody to detect GQ2. Kisspeptin-54 (Phoenix Pharmaceuticals Inc., Burlingame, Calif., USA) was used to preabsorb GQ2 for control purposes as described previously [21] (fig. 1).

E_2 -Containing Silastic Capsules

Crystalline E_2 (Steraloids Inc., Newport, R.I., USA) was used to prepare E_2 -filled Silastic capsules (4 cm long; Dow Corning Corp., Midland, Mich., USA) as described previously [22]. Blank Silastic capsules of the same dimensions were also prepared for the control experiments.

Nickel-Intensified Diaminobenzidine Immunohistochemistry

Sections were rinsed at room temperature in 50 mM potassium PBS (KPBS; pH 7.3, 6×10 min), incubated for 30 min in 3% hydrogen peroxide (Fisher Scientific, Fair Lawn, N.J., USA), and rinsed again in 50 mM KPBS (4×5 min). They were then incubated in blocking serum buffer [5% normal rabbit serum + 0.005% albumin from bovine serum (Sigma-Aldrich, St. Louis, Mo., USA) + 0.05% Triton X-100 (Sigma-Aldrich)] for 1 h to reduce nonspecific binding, and then incubated in GQ2 (1:60,000) for 1 h at room temperature followed by 48 h of incubation at 4°C on a shaker. After 10×6 min rinses in 50 mM KPBS, the sections were incubated in biotinylated rabbit anti-sheep IgG (1:600 in blocking serum) for 1 h. They were rinsed again in 50 mM KPBS (5×10 min) and incubated in a Vectastain Elite ABC kit (Vector Laboratories) for 1 h. This was followed by rinses in 50 mM KPBS and 0.175 M sodium acetate (3×5 min each). The tissue was then incubated in nickel-intensified diaminobenzidine (Ni-DAB) solution [nickel(II) sulfate hexahydrate (Sigma-Aldrich) and 3,3'-diaminobenzidine tetrahydrochloride hydrate (Sigma-Aldrich)]. The duration of incubation in the Ni-DAB, generally 3–8 min, was adjusted for optimal staining. To stop the reaction, the sections were rinsed with 0.175 M sodium acetate (3×5 min) followed by 3×5 min rinses in 50 mM KPBS. The sections were mounted on slides (Fisherbrand Superfrost Plus, Fisher Scientific, Pittsburgh, Pa., USA) treated with subbing solution (0.1% gelatin and 0.01% chromium potassium sulfate). The mounted sections were placed in distilled water for 2 min and then dehydrated in grades of ethyl alcohol (50, 75, and 100% twice for 4 min each). Before coverslipping with Permount (Fisher Scientific, Fair Lawn, N.J., USA), the slides were immersed in Histo-Clear (National Diagnostics, Atlanta, Ga., USA) for 4 min. The slides were stored at room temperature.

Cell Location and Cell Counting

The atlas of the monkey brain by Paxinos et al. [23] was used as a reference. Here, it should be noted that while this atlas identifies a specific anteroventral periventricular nucleus (AVPe) in the POA of the monkey brain, this is not the case for the atlas of Bleier [24]. The location of the AVPe in the POA of the monkey corresponds closely to that of the anteroventral periventricular area in rodents and sheep. For enumeration of the kisspeptin cell body

Fig. 2. Photomicrographs ($\times 40$) of hemisections of POA from a monkey showing neurons with 3 distinct patterns of kisspeptin immunostaining, giving the perikaryon a pear-like (a), granular (b), or cap-like (c) appearance. **Insets** Neurons demarcated by dashed boxes at higher magnification. Scale bars = 20 μm .

number, a series of coronal sections from individual monkeys taken every 250 μm from the level of the OVLT and continuing posteriorly to the caudal aspect of the optic chiasm were examined at $20\times$ magnification, and the neuron number per section was determined by blinded observers. The section containing the highest number of kisspeptin neurons for a given monkey was designated 'section 0', and the overall distribution of immunoreactive kisspeptin neurons for a given group of animals was derived by aligning the counts with respect to section 0, as previously described [25].

Hormone Assays

Plasma LH levels were measured as previously described, using a homologous macaque radioimmunoassay (RIA) employing recombinant cynomolgus monkey LH (rcLH; AFP6936A) as a standard [26]. The sensitivity of the assay ranged from 0.09 to 0.42 ng/ml, and the inter- and intra-assay coefficients of variation were less than 5 and 13%, respectively.

Plasma E_2 levels were measured using a Double Antibody RIA kit (Diagnostics Products Corp., Los Angeles, Calif., USA) as described previously [22]. The sensitivity of the assay ranged from 3 to 5 pg/ml, and the intra- and interassay coefficients of variation were less than 7 and 11%, respectively.

Plasma progesterone concentrations were measured using an RIA previously described [27]. The sensitivity of the assay ranged from 0.02 to 0.03 ng/ml, and the intra- and interassay coefficients of variation were less than 13 and 24%, respectively.

Statistical Analyses

The significance of differences in circulating E_2 and LH levels and in mean numbers of kisspeptin neurons per section were assessed using one-way ANOVA (Prism 5, GraphPad Software Inc., La Jolla, Calif., USA). Significance was accepted at $p \leq 0.05$.

Results

Experiment 1: Patterns of Kisspeptin Immunoreactivity in POA Perikarya and the Distribution of These Neurons in Intact Males and Females

Kisspeptin cell bodies in the POA of both male and female rhesus monkeys exhibited 3 distinct patterns of cytoplasmic kisspeptin immunoreactivity that gave the cell body the appearance of either (1) a homogeneously filled 'pear' generally with a pale central nucleus, (2) a granular and irregularly filled 'balloon', or (3) a homogeneous 'cap' typically located in the region of a dendrite (fig. 2). In the female, the predominant pattern was that of the homogeneous 'pear', while in the male, granular and irregular cytoplasmic staining prevailed. Kisspeptin perikarya in the POA of both males and females were distributed in an anteroposterior continuum spanning up to 2,750 μm (fig. 3, 4). For quantification, the 3 types of immunostained kisspeptin neuron were counted. Based on the topology of the base of the third ventricle, the greatest number of kisspeptin perikarya was found at an anteroposterior level corresponding to the region of the AVPe [23]. The number of immunoreactive kisspeptin cell bodies recorded in the analyzed sequence of sections through the POA of males and females was 591 ± 91 and 477 ± 79 (mean $\pm$ SEM), respectively. The difference was not significant. The distribution and intensity of immunoreactive kisspeptin fibers, within the region of the AVPe and beyond, was also similar in males and females (fig. 3).

Experiment 2: Immunoreactive Kisspeptin Expression in the POA Associated with an E₂-Induced LH Surge Prematurely Elicited in the Early Follicular Phase

Implantation of the E₂-containing Silastic capsules on day 3 of the control menstrual cycle in the 4 monkeys re-

sulted in an increase in circulating E₂ concentrations from a mean of 46 to a mean of 232 pg/ml (fig. 5a). Between 24 and 36 h after implantation of the E₂, an LH surge was elicited in all 4 monkeys, with peak LH concentrations being observed at 48 h (fig. 5a). In the 2 monkeys which subse-

Fig. 3. Photomicrographs ($\times 10$) of sequential hemisections taken at 250- μ m intervals through the POA of an intact adult male (**a-e**) and a female in the follicular phase of the menstrual cycle (**f-j**) stained for kisspeptin immunoreactivity. The sections are shown in order from anterior to posterior (male: **a-e**; female: **f-j**). Areas

demarcated by dashed boxes in the low-magnification panels are shown underneath at higher magnification ($\times 40$). SoR = Supra-optic recess of the third ventricle. Scale bars = 100 μ m (**a-j**, upper panels) and 20 μ m (smaller panels below **a-j**).

quently received E₂ during the experimental cycle, the increment in circulating E₂ was similar to that in the control cycle, and an LH surge was initiated at 24 h (fig. 5c). By the time of perfusion at 36 h, LH values were comparable to peak concentrations of this gonadotropin during the control cycle. As anticipated, implantation of blank Silastic capsules during the experimental cycle in the remaining 2 monkeys was not associated with an LH surge (fig. 5b).

Kisspeptin immunoreactivity in cell bodies in the region of the AVPe was markedly upregulated in the E₂-treated females when compared to the animals that received empty implants (fig. 6). Because there were only 2 animals in each group, kisspeptin cell numbers in this experiment were not enumerated.

Experiment 3: A Comparison of Immunoreactive Kisspeptin Neurons in the POA of Intact and Castrate Adult Male Monkeys and the Effect of E₂ Treatment on the Castrates

In this experiment, kisspeptin-immunoreactive perikarya and fibers in the POA were examined in 4 groups

Fig. 4. Mean number ($\pm$ SEM) of immunoreactive kisspeptin (KP-ir) neurons in the POA of intact males (open bars; $n = 3$) and females in the follicular phase (black bars; $n = 3$) enumerated in a series of sections taken at 250- μ m intervals through the POA. For each animal, the section containing the highest number of kisspeptin neurons was assigned section 0. Each data bar is composed of 1–3 observations.

Fig. 5. **a** Mean ($\pm$ SEM) concentrations of circulating E₂ and LH in females during a control menstrual cycle to demonstrate the ability of an E₂ challenge early in the follicular phase (day 3) to elicit a premature LH surge ($n = 4$). **b, c** Mean E₂ and LH levels in the 2 monkeys that were subsequently implanted with either E₂-containing capsules ($n = 2$; **b**) or blank capsules ($n = 2$; **c**). The arrows denote the time of implantation with either E₂-containing (E₂) or blank (BL) capsules.

Fig. 6. Photomicrographs ($\times 10$) of hemisections of POA at the level of the AVPe showing kisspeptin-immunoreactive neurons in intact females implanted on day 3 of the follicular phase with either blank (**a–d**) or E_2 -containing (**e–h**) Silastic capsules to induce a

premature LH surge. The areas demarcated by the dashed boxes in **a**, **c**, **e**, and **g** are shown at higher magnification ($\times 40$) in **b**, **d**, **f**, and **h**, respectively. SoR = Supraoptic recess of the third ventricle. Scale bars = 100 μm (**a**, **c**, **e**, **g**) and 20 μm (**b**, **d**, **f**, **h**).

of adult male monkeys, namely intact, castrate, and 2 cohorts of castrates exposed to approximately 300 pg/ml E_2 , one cohort for 12 h and the other for 48 h. Prior to conducting this study, the LH response to the E_2 signal was first characterized in all 6 E_2 -treated castrate monkeys (fig. 7a; note that the data in this panel are plotted with respect to the time of the LH peak, i.e. time 0). Implantation of the E_2 -containing Silastic capsules resulted in a marked increase in circulating E_2 from low castrate levels of <10 pg/ml to a mean concentration of approximately 1,000 pg/ml 12 h after implantation. Subsequently, the E_2 concentrations declined progressively, although they remained above 300 pg/ml for the duration of the time course (fig. 7a). Circulating LH concentrations declined from a mean of 6.9 ± 1.2 ng/ml before implantation of the

E_2 (fig. 7a) to 3 ± 1.1 ng/ml 12 h later. Subsequently, LH concentrations rose in all 6 animals, reaching peak levels of approximately 8 ng/ml 48–72 h after implantation of the E_2 -containing capsules (fig. 7a). During the subsequent 12- and 48-hour E_2 challenges, conducted 4–6 weeks after removing the capsules initially implanted for characterizing the LH response, circulating levels of this steroid were again increased from low castrate values to approximately 300 pg/ml, and, in both E_2 groups, circulating LH concentrations were reduced by approximately 50% after 12 h of steroid exposure (fig. 7b, c). In the 3 monkeys in which E_2 administration continued for 48 h, an unambiguous LH surge was observed in 1 animal. In this monkey, the LH concentrations increased 400% between 36 and 48 h.

Fig. 7. **a** Mean ($\pm$ SEM) circulating concentrations of E₂ and LH in castrate adult male monkeys ($n = 6$) implanted with E₂-containing capsules for 96 h during a control experiment to confirm the action of E₂ to induce an LH surge. **b, c** Mean E₂ and LH levels during a subsequent implantation of the same monkeys with E₂-containing capsules for either 12 h ($n = 3$; **b**) or 48 h ($n = 3$; **c**). Arrows denote the time of implantation with E₂-containing capsules (E₂). Time 0 represents the time of the LH peak (**a**) or the time of E₂ implantation (**b, c**). Pre = Prior to E₂ implantation. * $p \leq 0.05$ from Pre for E₂; ** $p \leq 0.05$ from -36 h for LH.

In striking contrast to the situation reported for the intact males in experiment 1, immunoreactive kisspeptin perikarya were very rarely seen in the POA of castrate adult male monkeys, although fiber staining in this region of the brain continued to be observed (fig. 8). In castrates treated with E₂ for either 12 or 48 h, however, the pattern and intensity of immunoreactive kisspeptin in perikarya of the POA closely resembled that seen in intact animals (fig. 8). In the E₂-treated castrates, kisspeptin cell bodies were observed along an anteroposterior continuum of 1,500 μ m in the region of the AVPe. The mean numbers of immunoreactive kisspeptin cell bodies in each of the 4 groups are shown in figure 9. E₂-treated castrates had a significantly greater number of kisspeptin neurons in the POA than had the castrate group, although the kisspeptin neuron number was unrelated to the duration of E₂ exposure.

Discussion

Although *KISS1*-expressing neurons have been previously described in the POA of both male and female monkeys [7, 8, 19], the present study is the first to confirm the expectation that, in these primates, *KISS1* mRNA in the POA is translated, and that kisspeptin may be readily detected in cell bodies and fibers in this hypothalamic area of both sexes by a standard immunohistochemical procedure. In addition, and more importantly, the present study provides evidence to indicate that the maintenance of the kisspeptin content in the POA in male primates is dependent upon the testis and that this action of the male gonad may be accounted for by E₂ receptor (ER) signaling.

As discussed in the Introduction, the POA in highly evolved primates is not essential for generating an LH surge and triggering ovulation [28]. Interestingly, however, Smith et al. [8] found that *KISS1* expression in the POA of the rhesus monkey is upregulated late in the follicular phase in association with an elevation in circulat-

Fig. 8. Representative photomicrographs ($\times 10$) of hemisections of POA at the level of the AVPe stained for kisspeptin immunoreactivity from intact (**a, e**), castrate (**b, f**), 12-hour E_2 -treated castrate (**c, g**), and 48-hour E_2 -treated castrate (**d, h**) adult male monkeys.

The high-power ($\times 20$) images below (**e-h**) show the areas demarcated by the dashed boxes in the top panels (**a-d**). SoR = Supra-optic recess of the third ventricle. Scale bars = 100 μm (**a-d**) and 50 μm (**e-h**).

ing E_2 concentrations that in primates, as well as in rodents, constitutes the ovarian signal that triggers the preovulatory LH surge [1, 29]. In the present study, imposition of an increment in circulating E_2 concentrations of approximately 300 pg/ml early in the follicular phase in 2 monkeys resulted in an unequivocal increase in the number of immunoreactive kisspeptin neurons in the region of the AVPe in association with the initiation of a premature LH surge. This result, albeit achieved with a limited number of animals, supports the earlier finding of Smith et al. [8] and reinforces the view that the population of kisspeptin neurons in the POA of the primate hypothalamus plays a role in mediating the positive feedback action of E_2 that elicits the preovulatory surge of LH in the normal menstrual cycle. The latter idea, however, must be tempered with the view – derived from the classic studies by Knobil's laboratory – that participation of the POA is

not obligatory for eliciting an LH surge and triggering ovulation in the monkey [reviewed in 28]. In order to fully appreciate the relative physiological importance of the populations of kisspeptin neurons of the POA and MBH in mediating the positive feedback action of E_2 on LH secretion in the monkey, selective ablation of ER signaling in these two populations of hypothalamic neurons will be required.

Since the neural mechanisms underlying the initiation of the preovulatory LH surge in the rhesus monkey are not decommissioned by the testis during perinatal development [14, 16], the finding that the number and distribution of kisspeptin neurons in the POA of testis-intact male monkeys was similar to that of female monkeys in the follicular phase of the menstrual cycle is perhaps not surprising. In both sexes, kisspeptin perikarya were observed in the region of the AVPe extending periventricu-

Fig. 9. Mean number ($\pm$ SEM) of immunoreactive kisspeptin (KP-ir) neurons in a sequential series of sections (8–11) collected at 250- μ m intervals throughout the POA of intact males, castrate males, and castrate males treated with E₂ for 12 or 48 h. ^{a, b} $p \leq 0.05$ from intact and castrate group, respectively.

larly into the medial POA, the same general area previously reported to contain *KISS1*-expressing neurons in the monkey [8] and other species [6]. There were, however, differences between the males and females in the distribution of immunoreactive kisspeptin within the perikarya of these neurons. In the males, there was a much larger proportion of neurons with cell bodies that appeared to be incompletely filled with immunoreactive kisspeptin. The reason for this subtle sex difference is unknown, but is likely to be related to the respective hormonal milieu generated by the testis or ovary. It should also be recalled that the method of fixation of the male (perfusion) and female (immersion) brain was different, although it is to be expected that perfusion rather than immersion would guarantee better preservation of immunoreactivity.

In striking contrast to intact males, immunoreactive kisspeptin perikarya were very rarely seen in the POA of adult males that had been castrated postpubertally. Implantation of castrate males for either 12 or 48 h with E₂-containing capsules that produced a quasi-square wave increment in circulating concentrations of the steroid of 300 pg/ml or greater resulted in a marked upregulation of kisspeptin immunoreactivity – an effect that was sufficiently robust to restore the number of immunoreactive kisspeptin neurons to values that were not significantly different from those in the POA of intact males. Here, it should be noted that the castrate group did not receive

blank Silastic capsules, nor were they sedated and venipunctured at 12-hour intervals before perfusion, as were the E₂-treated castrates. This minor flaw in the experimental design is considered to be of little importance, because the number of kisspeptin neurons and immunoreactive kisspeptin fibers in the groups of animals that were sedated and venipunctured (E₂ treated) was dramatically greater than that in the untreated castrate group. Intuitively, it would be anticipated that the effect, if any, of the additional experimental manipulations in the estrogen-treated groups would be to exert a suppressive influence on the kisspeptin neuronal network underlying gonadotropin secretion.

The action of E₂ treatment to regulate kisspeptin immunoreactivity in the POA of the castrate male monkey appeared to be unrelated to the action of the steroid to regulate LH secretion. This is because the kisspeptin immunoreactivity was dramatically upregulated at both 12 h, when LH secretion was suppressed due to the negative feedback action of E₂, and at 48 h, when the positive feedback action of the steroid was anticipated to be operative. The similarity in kisspeptin response at the two phases of E₂ feedback in the present model is consistent with the generally accepted notion that the site of action of E₂-negative feedback in primate and nonprimate species is within the MBH-pituitary unit [6, 29]. Indeed, E₂ replacement to the ovariectomized monkey results in the downregulation of both *KISS1* mRNA and kisspeptin in neurons of the arcuate nucleus in the MBH [30]. Thus, it is reasonable to propose that the kisspeptin content of kisspeptin neurons in the POA of the male monkey is maintained by an action of the testis, and that this action may be mimicked by high circulating concentrations of E₂ presumably via increasing signaling by ER α expressed by the kisspeptin neurons [31, 32]. The pharmacological levels of circulating E₂ achieved in the present study should not necessarily be used as an argument against the view that ER α signaling is the physiological pathway. This is because in the testis-intact situation, E₂ in the POA of the male will be generated primarily by the local aromatization of circulating testosterone [33, 34], which in the intact adult monkey ranges in concentration from 2,000 to 20,000 pg/ml [35]. Thus, the concentration of E₂ in the POA of intact males may be greater than that in the E₂-treated castrates examined in the present study.

In order for E₂ to exert its positive feedback action on LH secretion in the follicular phase of the menstrual cycle, elevated concentrations of the circulating steroid must be maintained for approximately 36 h [36]. If the strength-duration characteristics of the E₂-positive feedback signal

in the castrate male are similar to those in the intact female, the following question must be asked: what relevance, if any, does the upregulation of POA kisspeptin, observed in the present study after a 12-hour exposure to E_2 , have to do with the positive feedback action of the steroid and surge generation that is not anticipated until 24 h later? Clearly, to answer this question, there will be a need for a more discerning paradigm of E_2 replacement, such as that originally developed by Karsch et al. [15], wherein a 'pseudofollicular phase' is first induced in a castrate male by chronically increasing circulating E_2 to concentrations normally observed early in the follicular phase (i.e. 50 pg/ml) before imposing an additional elevation in E_2 to elicit positive feedback.

Two other issues need to be noted. First, kisspeptin staining in the untreated monkeys in the follicular phase of the menstrual cycle studied in experiments 1 and 2 appeared to be related to whether or not the animals had been implanted with blank Silastic capsules. In the implanted animals (experiment 2), the number of kisspeptin neurons was noticeably lower than that in the follicular-phase females without implants (experiment 1). A possible explanation for this discrepancy is that the monkeys which received Silastic capsules were sedated every 12 h – first to implant the capsules and then to collect bi-daily blood samples. Thus, the added stress imposed by the protocol employed in experiment 2 may have been responsible for the relatively low levels of kisspeptin immunoreactivity in this group of follicular-phase females.

Second, unambiguous evidence that the positive feedback action of E_2 had been initiated in the 3 castrate males that were exposed to E_2 for 48 h prior to perfusion was forthcoming in only 1 of the 3 monkeys. Although the circulating E_2 levels during the E_2 challenge at the time of perfusion were lower than those during the earlier pre-experimental challenge, when an LH surge was induced in all 3 of these monkeys, levels of the steroid were nevertheless maintained above 300 pg/ml. Therefore, an inadequate positive E_2 feedback signal is unlikely to account for variability of the LH response. A more likely explanation is that the LH surge had not yet been initiated in 2 animals at 48 h, a view consistent with the finding that during the pre-experimental E_2 challenge, the peak of the LH surge in the 3 monkeys was not observed until 60 or 72 h after implantation of the E_2 capsules.

In summary, the present study provides the first description of immunoreactive kisspeptin in the POA of the adult male and female macaque and suggests that the number and distribution of these peptidergic neurons is similar in the two sexes. In castrate adult males, however,

kisspeptin neurons were rarely observed in the POA, although kisspeptin immunoreactivity in this region of the brain was readily restored in male castrates following exposure to E_2 for as little as 12 h. The latter finding suggests that maintenance of the kisspeptin content in the POA of the intact male requires local synthesis of E_2 achieved via intrahypothalamic aromatization of circulating testicular testosterone. In addition, our results further support the view that kisspeptin neurons in the POA of the female monkey are a target for the positive feedback action of E_2 that underlies the initiation of the preovulatory LH surge, although this action of E_2 in highly evolved primates appears to be a redundant one and not obligatory for ovulation [4]. Finally, the presence of a robust kisspeptin neuronal network in the POA of the adult male monkey raises the question: what is the function of this peptidergic system in the brain of male primates?

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