Neurosteroids, trigger of the LH surge

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Abstract

Recent experiments from our laboratory are consistent with the idea that hypothalamic astrocytes are critical components of the central nervous system (CNS) mediated estrogen positive feedback mechanism. The “astrocrine hypothesis” maintains that ovarian estradiol rapidly increases free cytoplasmic calcium concentrations ([Ca\textsuperscript{2+}]\textsubscript{i}) that facilitate progesterone synthesis in astrocytes. This hypothalamic neuroprogesterone along with the elevated estrogen from the ovaries allows for the surge release of gonadotropin-releasing hormone (GnRH) that triggers the pituitary luteinizing hormone (LH) surge. A narrow range of estradiol stimulated progesterone production supports an “off-on-off” mechanism regulating the transition from estrogen negative feedback to estrogen positive feedback, and back again. The rapidity of the [Ca\textsuperscript{2+}]\textsubscript{i} response and progesterone synthesis support a non-genomic, membrane-initiated signaling mechanism. In hypothalamic astrocytes, membrane-associated estrogen receptors (mERs) signal through transactivation of the metabotropic glutamate receptor type 1a (mGluR1a), implying that astrocytic function is influenced by surrounding glutamatergic nerve terminals. Although other putative mERs, such as mER\textbeta, STX-activated mER-G\alpha\textsubscript{q}, and G protein-coupled receptor 30 (GPR30), are present and participate in membrane-mediated signaling, their influence in reproduction is still obscure since female reproduction be it estrogen positive feedback or lordosis behavior requires mER\textalpha. The astrocrine hypothesis is also consistent with the well-known sexual dimorphism of estrogen positive feedback. In rodents, only post-pubertal females exhibit this positive feedback. Hypothalamic astrocytes cultured from females, but not males, responded to estradiol by increasing progesterone synthesis. Estrogen autoregulates its own signaling by regulating levels of mER\textalpha in the plasma membrane of female astrocytes. In male astrocytes, the estradiol-induced increase in mER\textalpha was attenuated, suggesting that membrane-initiated estradiol signaling (MIES) would also be blunted. Indeed, estradiol induced [Ca\textsuperscript{2+}]\textsubscript{i} release in male astrocytes, but not to levels required to stimulate progesterone synthesis. Investigation of this sexual differentiation was performed using hypothalamic astrocytes from post-pubertal four core genotype (FCG) mice. In this model, genetic sex is uncoupled from gonadal sex. We demonstrated that animals that developed testes (XYM and XXM) lacked estrogen positive feedback, strongly suggesting that the sexual differentiation of progesterone synthesis is driven by the sex steroid environment during early development.

Keywords

Estrogen; Progesterone; LH surge; Positive feedback; Estrogen receptor; Astrocyte

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1. Role of neuroprogesterone in female reproduction

Ovulation is a critical event in mammalian female reproduction. In rodents and primates, maturing ovarian follicles synthesize and secrete estrogens. Circulating estrogen levels increase until they activate the hypothalamic-pituitary axis producing a surge release of luteinizing hormone (LH). This is the estrogen positive feedback that triggers ovulation. During positive feedback, the same estrogens that inhibited the hypothalamus and pituitary gland now stimulate these cells [1]. Many of the steps in the positive feedback cascade have been elucidated. In particular, rising estrogen levels induce the synthesis of hypothalamic progesterone receptors (PRs), which are required for the LH surge [2–6]. Specifically, Chappel and Levine demonstrated that both transcription and activation of PRs in the hypothalamus are obligatory events in the stimulation of the gonadotropin-releasing hormone (GnRH) and LH surges in estradiol-primed, ovariectomized (OVX) rats [7]. Studies with PR knockout mice in vivo demonstrated that PR-A in the hypothalamus, but not PR-B, mediates the LH surge and sexual receptivity in estrogen-primed female mice [8]. Therefore, not only is a pre-ovulatory increase in peripheral estradiol required, but an increase in progesterone synthesis and activation of PRs are all essential for inducing the LH surge.

In the intact rat, both the ovary and the adrenal cortex are highly steroidogenic organs capable of producing the pre-ovulatory rise in progesterone needed for the LH surge [9]. However, no significant rise in progesterone has been detected in the systemic circulation prior to the LH surge, indicating that the progesterone required for the LH surge may not be synthesized peripherally [10–12]. Consistent with this idea is that neither the adrenals nor the ovaries are necessary for an estrogen-induced LH surge [13,14]. Indeed, OVX and adrenalectomized (ADX) rats injected with 17β-estradiol, but not progesterone, have been shown to produce a robust LH surge [13,15].

The source of this progesterone appears to be from the brain. The steroidogenic capacity of the brain has been well established [16–26]. Neuroprogesterone, progesterone synthesized de novo in the brain, can be induced by estradiol. Neurons, astrocytes, and oligodendrocytes have been demonstrated to possess all the steroidogenic enzymes and associated proteins required to convert cholesterol directly to progesterone within the brain, including cytochrome P-450 side-chain cleavage (P450scc), 3β-hydroxysteroid dehydrogenase (3β-HSD), steroidogenic acute regulatory protein (StAR), and 18 kDa translocator protein (TSPO), formerly known as peripheral-type benzodiazepine receptor (PTBR) [27,28]. Estradiol treatment of OVX/ADX female rats increased hypothalamic neuroprogesterone levels and induce a physiological relevant LH surge, indicating that the source of progesterone was neither the ovary nor adrenal gland [15]. Furthermore, treatment with trilostane, a blocker of the enzyme 3β-HSD that catalyzes the conversion of pregnenolone to progesterone, inhibited the LH surge, indicating that neuroprogesterone synthesis is critical for estrogen positive feedback in OVX/ADX female rats [15]. In gonadally intact rats with normal four-day estrous cycles, blocking hypothalamic steroidogenesis with aminoglutethemide (AGT), a P450scc enzyme inhibitor, on the morning of proestrus eliminated the LH surge, ovulation and luteinization [29]. After several days, the effects of AGT wore off, hypothalamic progesterone synthesis recovered, and the treated rats resumed their estrous cycles. These data strongly suggest that estrogen stimulates neuroprogesterone synthesis locally within the hypothalamus, which is essential (directly or through its metabolites) in mediating the positive feedback regulation of the LH surge.
2. Estrogen effects on astrocytes

Our understanding of astrocytes in regulating nervous system function has evolved from providing structural support to regulating metabolic events [30] and synaptic function in adjacent neurons [31,32]. Astrocytes respond to numerous transmitters, peptides, and steroids [33–36]. It is now well accepted that estradiol acts on astrocytes [37]. Similar to granulosa cells of the ovary, astrocytes have been shown to express estrogen receptor-alpha (ERα) and estrogen receptor-beta (ERβ), which provides a mechanism for estradiol regulation [33,38–42]. Estradiol profoundly influences astrocyte morphology and function [43], glial fibrillary acidic protein (GFAP) distribution [44], sexual differentiation [45], and steroidogenesis [46,47]. The presence of surrounding neurons further enhances the changes in astrocytic shape induced by estradiol [48]. Astrocytes have been described to play an important role in estrogen-mediated neuroprotection [49]. Astrocytes also regulate numerous hypothalamic processes including regulation of releasing factors [50–54] and synthesis of neurosteroids [54–57].

Although enriched cultures of neurons and oligodendrocytes are capable of synthesizing progesterone, enzymatic activity studies indicate that astrocytes are the most steroidogenically active cells in the brain [57]. Not only do astrocytes contain ERs and interact with neurons in response to estradiol, but astrocytes are the main source of the essential neuroprogesterone produced within the hypothalamus [15,37,46,57]. Thus, hypothalamic astrocytes are critical for the central nervous system (CNS) response mediating estrogen positive feedback [58,59]. This increase in hypothalamic neuroprogesterone activates the progesterone receptors in the neuronal circuit that regulates the activity of GnRH neurons, resulting in greater release of GnRH that triggers the pituitary LH surge leading to ovulation – the critical event in female reproduction [6,7,15,29,37].

3. mER signaling

As in neurons, estradiol can influence cell signaling in astrocytes, which express ERα and ERβ both intracellularly and in the plasma membrane [33,40–42]. Therefore, estradiol can activate nuclear-initiated and/or membrane-initiated signaling mechanisms. Classic nuclear-initiated estradiol action is well established and mediated through activation of ERα and ERβ located in the nucleus to behave as ligand-activated transcription factors. Evidence suggests that these same receptors can mediate both nuclear- and membrane-initiated signaling. Although long-studied, it is only more recently that membrane-initiated estradiol action has been widely accepted [33,42,60–76].

Activation of membrane-associated estrogen receptors (mERs) with estradiol or a membrane impermeable construct E-6–BSA (estradiol-coupled to bovine serum albumin) initiates a rapid [Ca^{2+}℄ increase via activation of the phospholipase C/inositol trisphosphate (PLC/IP3) pathway that releases intracellular stores of calcium from the smooth endoplasmic reticulum in neurons and astrocytes [33,46,77]. This rise in [Ca^{2+}℄ stimulates the de novo synthesis of progesterone in post-pubertal female hypothalamic astrocytes within 5 min [15,46,78]. Confirmation of this idea was obtained through the use of thapsigargin, a potent Ca^{2+} ATPase inhibitor that rapidly releases IP3-sensitive Ca2+ stores from the smooth endoplasmic reticulum. This massive release of Ca^{2+}, which was similar in magnitude to estradiol stimulation, resulted in progesterone synthesis by itself [46]. Although these studies were done with nanomolar doses of estradiol, subsequent experiments demonstrated that subnanomolar doses of estradiol were sufficient to induce [Ca^{2+}℄ release in cultured hypothalamic astrocytes [62]. Estradiol induction of progesterone synthesis had a half-maximal effector concentration (EC50) of 0.82 nM, which may be related to the extent of the [Ca^{2+}℄ increase [78]. Thus, the estradiol facilitation of progesterone synthesis appears to be
a “step function” responding to physiological levels of estradiol that are reached during the proestrus surge [79–81]. Both estradiol and progesterone stimulation of the hypothalamus are essential for estrogen positive feedback, ultimately leading to the LH surge [7,15,37,59]. The threshold response to estradiol is consistent with the idea that stimulation of neuroprogesterone synthesis is part of an “off-on-off” mechanism regulating the transition from estrogen negative feedback to estrogen positive feedback, and back again [78]. For example, as estradiol rises with developing ovarian follicles, gradually increasing levels of \([\text{Ca}^{2+}]_i\) release will be stimulated by the estradiol. However, only with physiologically peak estradiol levels, consistent with mature follicles ready for ovulation, does the \([\text{Ca}^{2+}]_i\) release reach a critical threshold allowing for progesterone synthesis. Otherwise, hypothalamic progesterone may rise too early, resulting in a premature LH surge before ovarian follicles are fully mature and ready to ovulate.

Membrane-impermeable E-6–BSA–FITC (estradiol–bovine serum albumin–fluorescein isothiocyanate conjugate) and E-6–biotin (estradiol–biotin conjugate) constructs also bind to and demonstrate membrane ERs as well as their internalization in neurons and astrocytes [58,82,83]. The estradiol effects were blocked by the ER inhibitor ICI 182,780, confirming non-genomic estradiol signaling through a mER [33,46]; however, how a membrane-associated nuclear receptor activates intracellular signaling cascades was not immediately clear. Membrane-initiated estradiol signaling (MIES) has been shown to activate G protein-dependent cell signaling cascades [84], including activation of the mitogen-activated protein kinase (MAPK) pathway, increasing \([\text{Ca}^{2+}]_i\), activation of protein kinase A (PKA) and protein kinase C (PKC), and phosphorylation of cAMP-responsive element binding protein (CREB) [85,86] reviewed by [87]. A mechanism for MIES was proposed in which ligand-bound mER\(\alpha\) or mER\(\beta\) transactivated metabotropic glutamate receptors (mGluRs) to stimulate PLC/IP\(3\)–MAPK pathways leading to the activation of CREB [85,88]. These membrane-initiated signaling cascades have been implicated in the estradiol activation of lordosis behavior through receptor activation and neuropeptide expression [89]. For steroid synthesis, activation of PKA and/or PKC is required for the phosphorylation and activation of StAR, the rate-limiting step in steroid biosynthesis [90–92]. In mammalian astrocytes, TSPO is another possible regulator of intramitochondrial cholesterol transfer [28,93]. Activation of the TSPO in whole animals increased brain levels of allopregnanolone, pregnenolone, and progesterone [94]. In astrocytes, TSPO agonists stimulated the synthesis of pregnenolone, the immediate precursor of progesterone [95].

In hypothalamic astrocytes, both the estradiol-induced \([\text{Ca}^{2+}]_i\) release and progesterone synthesis are blocked by the metabotropic glutamate receptor type 1a (mGluR1a) antagonist LY 367385 [62,78]. Co-immunoprecipitation of ER\(\alpha\) and mGluR1a is consistent with a mER–mGluR interaction. We observed an interesting phenomenon. Even though estradiol would transactivate the mGluR1a in the absence of glutamate, if the mGluR1a was selectively activated with DHPG, a mGluR1a agonist, the estradiol response was augmented (Fig. 1) [62,78]. High dose DHPG can be used to accurately predict the maximal effect on astrocytes from the glutamate released by surrounding neurons that would be present in vivo. Furthermore, surface biotinylation studies in hypothalamic astrocytes demonstrated that mER\(\alpha\) trafficking is dependant upon the mGluR1a, such that mGluR1a antagonism with LY 367385 blocked estradiol-dependant mER\(\alpha\) insertion and its internalization [60]. In hypothalamic neurons, estradiol treatment significantly increased the internalization of mGluR1a in parallel with ER\(\alpha\), further supporting an mER\(\alpha\)-mGluR1a signaling unit [96]. Thus, trafficking and internalization of mER\(\alpha\) likely occurs together as a complex with mGluR1a. These studies also suggested that membrane-initiated ER signaling may be required for the initiation of mER\(\alpha\)-mGluR1a trafficking to the membrane. The membrane impermeable E-6–BSA construct demonstrated increased mER\(\alpha\) trafficking similar to estradiol [97]. Internalization, a measure of receptor activation, closely mimicked the
mERα-mGluR1α trafficking. As expected, the more receptor complexes on the membrane, the more activation and internalization. In astrocytes, this is supported by the augmentation of the [Ca\(^{2+}\)]\(_i\) response during maximal mERα insertion and internalization at 30–60 min of estradiol exposure [60]. Overall, maximal signaling in hypothalamic astrocytes requires the presence of both glutamate and estradiol, implying that estradiol signaling is augmented by excitatory neural activity involving glutamate and astrocytes are a site of neural–hormonal integration (Fig. 1). Indeed, middle-aged female rats exhibit reduced excitation of GnRH neurons and attenuated LH surges compared to young females under estrogen positive feedback conditions, in part, due to decreased glutamate neurotransmission in the medial preoptic area, demonstrating the importance of local modulatory effects on estrogen positive feedback, the LH surge, and reproductive function [98,99].

4. One ligand, many receptors

In terms of cell signaling, the classic nuclear receptors ERα and ERβ have been shown to associate with the plasma membrane with a variety of methods. Overexpression of ERα and ERβ demonstrated that a percentage of the nuclear proteins are targeted to the plasma membrane [73] where they have been localized with immunohistochemistry, western blotting, and surface biotinylation [33,42,60–63,65,68,70,71,73,74,82,96,100,101]. In terms of reproduction, ER knockout mice in vivo demonstrated that ERα, but not ERβ, was critical for estrogen positive feedback and the LH surge [102]. In our laboratory, OVX ERα knockout mice (ERKO) mice in vivo treated with 17β-estradiol benzoate (10 μg) failed to significantly increase hypothalamic progesterone levels that were seen in OVX wild type mice [78]. Similar results were seen with in vitro studies using ERKO mouse astrocytes cultures such that the estradiol-induced [Ca\(^{2+}\)]\(_i\) response was significantly attenuated [78]. In addition, several other membrane-associated ERs that mediate rapid estrogen effects have been reported: ER-X [103,104], STX-activated protein called mER-Gq [69,105] and G protein-coupled receptor 30 (GPR30) [66,67,72,106].

We examined several of these putative mERs in our astrocrine model in which estradiol elicits an increase in [Ca\(^{2+}\)]\(_i\) stimulating progesterone synthesis [78]. Since we have demonstrated both ERα and ERβ in the plasma membrane fractions of astrocytes [33], we selectively activated them with PPT, an ERα agonist [107], and DPN, an ERβ agonist [108,109]. As expected, PPT mimicked the estradiol actions and was antagonized by the mGluR1α antagonist LY367385 [78]. Conversely, equimolar DPN did not increase [Ca\(^{2+}\)]\(_i\) nor progesterone synthesis in our hypothalamic astrocyte cultures [78]. Furthermore, mERα co-immunoprecipitated with mGluR1α indicating a potential interaction, but mERβ did not co-immunoprecipitate with mGluR1α [62,78]. These results support the idea that, of the two classic ERs, ERα appears to be the mER mediating estradiol actions for progesterone synthesis in hypothalamic astrocytes, consistent with estrogen receptor knockout studies demonstrating that ERα, not ERβ, is essential for estrogen positive feedback and the LH surge [102].

The location of mERs had been a subject of controversy with possible cytoplasmic localization associated with the inner phospholipid bilayer. However, surface biotinylation with a membrane impermeable reagent tags proteins that protrude from the plasma membrane and has recently been used to confirm the status of ERα as a membrane protein with an extracellular portion, possibly containing the ligand binding domain in both hypothalamic astrocytes and neurons [60,96,110]. Hydropathy analysis of ERα suggests a potential transmembrane domain near the amino-terminal domain (SOSUI, TMpred program), which support a potential extracellular ERα binding site. Unfortunately, the biotinylation experiments do not indicate which part of the ERα extends through the membrane. Interestingly, a 52–55 kDa ERα protein was also labeled by surface
biotinylation. This protein is an alternatively spliced ERα that is missing exon 4 (ERαΔ4) [97], which has been reported in the brain and breast [111–114]. Coincidentally, exon 4 of ERα codes for the hinge region directing nuclear localization, which may explain the preferential trafficking of ERαΔ4 to the plasma membrane. However, only the full-length 66 kDa ERα co-immunoprecipitated with mGluR1a, which in astrocytes is needed for the estradiol-induced [Ca^{2+}]_i release and progesterone synthesis [62,78].

A significant observation from these experiments was the autoregulation of estradiol signaling. ERα trafficking to and from the plasma membrane was rapid, such that 5 min of estradiol treatment significantly increased mERα insertion and internalization, suggesting that de novo synthesis of ERα is probably not occurring [60]. Insertion is most likely through estradiol-induced exocytosis of vesicles containing mERα. Such ERα-immunoreactive vesicles have been observed in hippocampal neurons [115,116] and pituitary cells [117]. Internalization, a common mechanism for regulating membrane signaling, is usually through the formation of endocytic vesicles [83]. Once receptors are internalized and release their ligand, they can either be recycled to the cell surface or degraded. This mERα trafficking and internalization is blocked by the ER antagonist ICI 182,780 and dependant upon the presence of estradiol, with estradiol-induced [Ca^{2+}]_i response correlating with maximal insertion and internalization of mERα [60]. Continuous exposure to estradiol eventually reduced levels of mERα and its internalization to basal levels, suggesting a down-regulation of the receptor that temporally limits membrane-initiated cell signaling [60,96]. Membrane levels of ERα remained low for 24–48 h after estradiol exposure [60]. At some point after this down-regulation, mERα levels are partially restored. Currently, the time course for recovery of mERα levels is not known. If mERα-mGluR1a is a signaling unit, then blockade of the mGluR1a should prevent both the insertion of ERα into the membrane and its internalization. This is exactly what was observed [60]. Therefore, levels of mERα are autoregulated by the concentration of estradiol in the surrounding extracellular environment, which determines the magnitude of the MIES response and its duration.

In addition to mERα and mERβ, several other candidate mERs have been proposed [69,72,103]. One putative mER is GPR30, a G protein-coupled receptor (GPCR) that activates adenylyl cyclase in breast cancer cells lacking both ERα and ERβ [66,67,72,106]. Although FLAG- and hemagglutinin-tagged GPR30 have been reported at the plasma membrane [118,119], GPR30 could not be identified at the plasma membrane or labeled with surface biotinylation in native cells [60,78,110]. The GPR30 agonist G-1, a substituted dihydroquinoline [120], stimulated [Ca^{2+}]_i release and progesterone synthesis [78]. However, G-1 seemed to signal through a different mechanism compared to estradiol since G-1 was not blocked by antagonizing the mGluR1a [78]. The lack of interaction between GPR30 and mGluR1a was confirmed by the absence of co-immunoprecipitation between these proteins [78]. While it is difficult to understand the discrepancy with GPR30 localization in the plasma membrane, our results can be interpreted to support the observation of estradiol activation of intracellular GPR30 on the endoplasmic reticulum [67]. Activation of GPR30 may directly induce the release of intracellular stores of Ca^{2+}, which in turn stimulates progesterone synthesis. However, only very large molar concentrations of G-1 activated [Ca^{2+}]_i release and progesterone synthesis [78]. Such a response was reminiscent of DPN, an ERβ agonist, which is not thought to be physiologically important in the regulation of the LH surge.

Another candidate receptor is a membrane-associated binding protein that is Gαq-coupled and activated by estrogen as well as STX, a diphenylacrylamide selective estrogen receptor modulator (SERM) [69]. STX remains efficacious in double ERα/ERβ knockout mice, but blocked with the ER antagonist ICI 182,780 [105]. STX does not activate ERα or ERβ.
having a million-fold lower binding affinity compared with estradiol for these receptors [69]. However, this STX-activated mER-Gαq also activates PLC [69], a signaling pathway similar to that activated by mERα-mGluR1a. In hypothalamic astrocytes, STX increased [Ca2+]i and progesterone synthesis through transactivation of mGluR1a [78]. It has been suggested that STX signals through GPR30, such that small interfering RNA directed against GPR30 abolished the STX-induced transcription [121]. However, estradiol has been proposed to signal through the STX-activated mER-Gαq pathway in GPR30 knockout mice [122]. In our hands, STX and G-1 produced distinctly different responses in hypothalamic astrocytes [78]. Although estradiol and STX responses are blocked by mGluR1a antagonism and activate the same PLC pathway, these actions are mediated through different receptors since STX stimulation of Ca2+ release remains in astrocytes from ERKO mice, where estradiol was ineffective [78]. Since ERKO mice do not demonstrate estrogen positive feedback and lack sexual receptivity due to the lack of estradiol effect, the STX response in ERKO mice is not consistent with a STX-related signaling mechanism for induction of the LH surge or receptivity in female reproduction [78,102,123,124]. Furthermore, this mER-Gαq has much lower affinity for estradiol (~20-fold) compared with STX [105]. Although STX was extremely potent compared with estradiol at stimulating the mER-Gαq in ERKO mouse astrocytes, STX does not exist in vivo and mER-Gαq activation by physiological estradiol levels seems too weak for any measurable signaling actions in parameters of reproduction. Therefore, the physiological relevance of the STX-activated mER-Gαq in reproductive function remains to be elucidated.

Lastly, ER-X has been proposed as an estrogen receptor during development and following injury, especially in the cortex [103,104]. This mER is not inhibited by ICI 182,780, but is activated by 17β-estradiol. Interestingly, ER-X is unique in that it is preferentially activated by 17α-estradiol [104]. However, in astrocytes, the estradiol action on [Ca2+]i is stereospecific [33] and the 17β-estradiol-induced [Ca2+]i release and progesterone synthesis was inhibited by ICI 182,780, which are not consistent with an ER-X mediated action [46,62].

In summary, several putative mERs are involved in regulation of [Ca2+]i release and progesterone synthesis in hypothalamic astrocytes [78]. The evidence was strongest for ERα and the STX-activated mER-Gαq, both of which required transactivation of mGluR1α to initiate cell signaling. From a reproductive vantage point as well as evidence from wild-type and ERKO hypothalamic astrocytes, ERα is the primary mER responsible for the rapid cell signaling that leads to an increase in hypothalamic neuroprogesterone [46,60,62,78].

5. Sex differences in estradiol stimulated progesterone release

Although both male and female rodents have a well-developed negative feedback mechanism regulating the release of GnRH and LH, one of the most robust sexually differentiated physiological responses is estrogen positive feedback, which induces the surge release of LH in response to estradiol stimulation. This phenomenon of estrogen positive feedback is a hallmark of various female animal species. Males, especially male rodents, do not exhibit this phenomenon. For rodents, once the ability to produce estrogen positive feedback is lost during development, this loss is permanent. In primates, including humans, many years of continuous estrogen exposure in males can result in an estrogen positive feedback response, although it is quite attenuated [125]. According to the epigenetic theory of sexual differentiation of the brain, the sex difference in estrogen positive feedback arises from the action of estradiol (aromatized from testosterone) during organization of the neural circuit(s) controlling the GnRH neurons [126–131]. Several mechanisms have been proposed to account for this differentiation, including the lack of estrogen-induced synaptic plasticity in the male arcuate nucleus [132] and an attenuated distribution of kisspeptin...
neurons in males [133]. Various structural sex differences that result from perinatal exposure to estradiol have been identified. In terms of regulating GnRH, males have greater postnatal apoptosis in the developing anteroventral periventricular nucleus (AVPV), a region crucial for estrogen positive feedback in females [134–136]. Although it is not clear whether such a sex difference in apoptosis is an important mechanism for elimination of estrogen positive feedback, it does support a role for postnatal sex steroids in organizing brain mechanisms involved in reproduction [137].

A mechanism for mediating estrogen positive feedback, the LH surge, and transition from proestrus to estrus involves the synthesis of neuroprogesterone by astrocytes in the hypothalamus [58,59]. Significantly, estradiol treatment in vivo increased progesterone levels in the female, but not male, hypothalamus [15,58]. In other words, males and reproductive senescent females, which do not have increased hypothalamic progesterone synthesis in response to estradiol, fail to demonstrate an estrogen positive feedback mechanism. Investigation of astrocytic sex differences in the post-pubertal hypothalamus of rodents further confirmed astrocytes and neuroprogesterone as a critical feature in the neurosteroid regulation of the LH surge. Male astrocytes had a significantly attenuated estradiol-induced $[\text{Ca}^{2+}]_i$ response and failed to synthesize progesterone in contrast to female astrocytes (Fig. 2) [37,46,78,138]. This was consistent with previous observations that only post-pubertal female rodents have increased levels of progesterone in the hypothalamus before the LH surge [15]. There appears to be a crucial concentration of $[\text{Ca}^{2+}]_i$ required for neuroprogesteone synthesis. For example, male astrocytes had an attenuated $\text{Ca}^{2+}$ response that was unable to facilitate progesterone synthesis [138], which is consistent with previous reports in neonatal astrocytes [46] and in post-pubertal astrocytes, in which 0.1 nM estradiol stimulated $[\text{Ca}^{2+}]_i$ release [62], but not progesterone synthesis [78]. One possible mechanism that could produce the decreased $[\text{Ca}^{2+}]_i$ response is the attenuated insertion of $\text{mER}_\alpha$ into male astrocytes compared with female astrocytes [138].

Biological differences between males and females can result genetically from direct sex chromosome differences, developmentally through differential exposure to sex steroids during developmental “organization”, or functionally from acute “activational” effects of gonadal steroids operating at many life stages, which can be controlled through gonadectomy [139]. Perinatal gonadal hormone secretions have been shown to have powerful and permanent actions on physiology, including pituitary function, gene expression in the brain and sexual behavior [140–143]. In spite of these epigenetic effects, several chromosomal dependent sex differences have been demonstrated in the brain. Specifically, the four core genotype (FCG) mice model has demonstrated purely chromosomal XX versus XY differences in behaviors, including aggression, parenting, habit formation, nociception and social interactions [144]. In addition, $\text{Sry}$ is expressed in the brain, and it has been shown to directly influence the biochemical properties of the dopaminergic neurons of the nigrostriatal system and the specific motor behaviors they control [145]. Although a stark difference between male and female astrocytic response to estradiol was demonstrated, it was not clear whether the differences result from the sex chromosome complement or to the presence of the $\text{Sry}$ gene with its influence on gonadal development and the early sex steroid environment. Using FCG mice, animals with ovaries (XXF and XYF) had astrocytes in which estradiol facilitated progesterone synthesis, regardless of whether they had one or two X chromosomes [138]. Conversely, mice with testes (XYM and XXM) were unresponsive to estradiol and did not increase progesterone synthesis [138]. These results suggest a $\text{Sry}$ transgene effect and not a sex chromosome effect on hypothalamic astrocyte response to estradiol. The effects from the $\text{Sry}$ transgene could be due to direct effects of the $\text{Sry}$ gene itself or its influence on gonadal differentiation and the sex steroid environment during early development. Interestingly, the XYM from FCG mice synthesized little or no progesterone [138], which could suggest a potential
chromosomal effect. However, male wild type astrocytes, without Sry translocation to an autosome, synthesized basal progesterone levels similar to female wild type astrocytes [138]. Therefore, this difference could potentially be caused by the deletion and transgenic insertion of Sry, resulting in the inactivation of surrounding gene(s), positional effects or differential expression of the Sry transgene. Differences between wild type XY males and FCG XYM have been previously reported for mounting behavior, social exploration and concentration of tyrosine hydroxylase-immunoreactive neurons within the AVPV [146]. Unfortunately, the steroid profile of XYM in FCG mice has not yet been characterized.

6. Conclusion

Membrane ERs are clearly important for signaling in astrocytes as well as in neurons. Our studies have addressed several key questions: can ERα be a membrane receptor, how does it signal, what regulates its signaling, and is it sexually differentiated. Estradiol binds to mERα to transactivate mGluR1a, which activates the PLC/IP3 pathway leading to the release of IP3 receptor-sensitive stores of intracellular Ca2+. The rapid increase in [Ca2+]i activates various kinases resulting in the phosphorylation of enzymes that facilitate neuroprogesterone synthesis essential for estrogen positive feedback, the LH surge, and ovulation. In contrast to female astrocytes, male astrocytes had an attenuated [Ca2+]i release that was unable to facilitate progesterone synthesis. One possible mechanism for this sexually differentiated response is the attenuated trafficking of mERα to the plasma membrane in male astrocytes that may result from differential sex steroid exposure during early development.

We have also demonstrated that other putative ERs are present in astrocytes and participate in cell signaling, but their influence on reproductive function is still obscure. Female reproduction be it lordosis behavior or estrogen positive feedback requires ERα. In hypothalamic astrocytes, high doses of DPN were required to induced estradiol-like [Ca2+]i responses, most probably through weak non-selective mERα activation rather than mERβ activation, but even high doses of DPN failed to stimulate progesterone synthesis. Our studies do not resolve the role of GPR30 in MIES. We and others have not been able to demonstrate native GPR30 in the plasma membrane. GPR30 may uniquely mediate estradiol-like responses exclusively through an intracellular receptor on the endoplasmic reticulum. This is supported by the high concentration of G-1 required to stimulate a [Ca2+]i response and progesterone synthesis. Its dose response resembles that of DPN, which does not seem to play a critical role in the regulation of estrogen positive feedback.

Unfortunately, the STX-activated mER-Gαq has yet to be identified. STX was extremely potent compared with estradiol at stimulating the mER-Gαq in ERKO mouse astrocytes, but physiological estradiol levels seem insufficient to stimulate any significant reproductive signaling effects through this receptor. To fully elucidate the role of GPR30 and the STX-activated mER-Gαq, additional experiments will be required. Overall, mERα in post-pubertal female hypothalamic astrocytes seems to play an essential role in sensing the rising levels of estradiol during proestrus to regulate neuroprogesterone levels critical for regulating estrogen positive feedback and the LH surge.

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Fig. 1.
An illustration of the additive effects of glutamate released from surrounding neurons on the estradiol response in hypothalamic astrocytes. Estradiol acting upon mERα induces $[Ca^{2+}]_i$ release that stimulates progesterone synthesis essential for estrogen positive feedback and the LH surge. Local neuronal activity modulates hypothalamic astrocyte function through the release of glutamate, which activates the mGluR1a to augment both the estradiol-induced $[Ca^{2+}]_i$ response and progesterone synthesis. Only when estradiol bound mERα interacts with glutamate bound mGluR1a does maximal intracellular signaling take place in hypothalamic astrocytes, suggesting that astrocytes are a site of neural–hormonal integration.
Fig. 2.
Sex differences in estradiol stimulated progesterone synthesis in post-pubertal hypothalamic astrocytes. Estradiol at 1 nM and 100 nM stimulated significant progesterone synthesis in female astrocytes ($p < 0.05$ vs. control). Conversely, male astrocytes did not synthesize progesterone ($p > 0.05$ vs. controls) when stimulated by estradiol at 1 nM or 100 nM.
*Significantly different compared to female control and all male groups ($p < 0.05$, one-way ANOVA with Student–Newman–Keuls post hoc test).
†Significantly different compared to female astrocytes stimulated with 1 nM estradiol ($p < 0.05$, one-way ANOVA with Student–Newman–Keuls post hoc test).

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