The Role of Kiss1 Neurons As Integrators of Endocrine, Metabolic, and Environmental Factors in the Hypothalamic–Pituitary–Gonadal Axis

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Kisspeptin–GPR54 signaling in the hypothalamus is required for reproduction and fertility in mammals. Kiss1 neurons are key regulators of gonadotropin-releasing hormone (GnRH) release and modulation of the hypothalamic–pituitary–gonadal (HPG) axis. Arcuate Kiss1 neurons project to GnRH nerve terminals in the median eminence, orchestrating the pulsatile secretion of luteinizing hormone (LH) through the intricate interaction between GnRH pulse frequency and the pituitary gonadotrophs. Arcuate Kiss1 neurons, also known as KNDy neurons in rodents and ruminants because of their co-expression of neurokinin B and dynorphin represent an ideal hub to receive afferent inputs from other brain regions in response to physiological and environmental changes, which can regulate the HPG axis. This review will focus on studies performed primarily in rodent and ruminant species to explore potential afferent inputs to Kiss1 neurons with emphasis on the arcuate region but also considering the rostral periventricular region of the third ventricle (RP3V). Specifically, we will discuss how these inputs can be modulated by hormonal, metabolic, and environmental factors to control gonadotropin secretion and fertility. We also summarize the methods and techniques that can be used to study functional inputs into Kiss1 neurons.

Keywords: Kiss1, hypothalamus, arcuate nucleus, gonadotropin-releasing hormone, neural afferents, luteinizing hormone

INTRODUCTION

Kisspeptin, encoded by the Kiss1 gene, was initially proposed as a suppressor of metastasis, but its precise role in this process remains elusive (1, 2). Expression of the KISS1 gene and the gene encoding its cognate receptor, the G-protein coupled receptor 54 (GPR54, also called KISS1R) has been detected in the brain and peripheral tissues, including the pancreas, liver, small intestine, pituitary, and placenta (1, 3, 4). Two seminal studies in 2003 defined a physiological role for kisspeptin by showing that inactivating mutations of the GPR54 gene are associated with hypogonadotropic hypogonadism in humans and mice (5, 6). Kisspeptin signaling was thus identified as one of the critical regulators for both puberty onset and maintenance of normal reproductive functions in mammals (7, 8). Kisspeptin exerts its effects on the hypothalamic–pituitary–gonadal (HPG) axis, by acting as a neupeptide essential for stimulation of gonadotropin-releasing hormone (GnRH) neurons in the hypothalamus (9–13). Subsequently, kisspeptin signaling has also been implicated in regulating glucose homeostasis and body weight control (14, 15).

This review focuses mainly on how Kiss1 neurons in the arcuate region of the hypothalamus (Kiss1ARC) act as sensors to relay information about hormonal and nutritional status and environmental changes to GnRH neurons to regulate the secretion of the gonadotropins critical to sustain fertility (Table 1). It is important to decipher the upstream networks that connect to Kiss1ARC.
Table 1: Summary of neuropeptides or hormones that interact with Kiss1 neurons.

| Candidates                      | Anatomical evidence to ARC or Kiss1 neurons | ICV or electrophysiology studies | Receptors expressed in Kiss1 neurons | Physiological relevance |
|---------------------------------|--------------------------------------------|----------------------------------|--------------------------------------|-------------------------|
| Agouti-related peptide/neuropeptide Y (AgRP/NPY) | AgRP neurons formed synaptic connections with Kiss1ARC neurons (mice) (135) | Channel rhodopsin-assisted mapping indicated AgRP neurons formed inhibitory synaptic connections with Kiss1ARC neurons (mice) (135) | Npy1r, Npy2r, and Npy5r (82) | Chemogenic activation of AgRP neurons inhibited fertility in vivo (135) |
| Arginine vasopressin (AVP)      | AVP-immunoreactive fibers in close apposition with Kiss1impeu neuron (mice) (32) | AVP-stimulated Kiss1impeu neurons (105) | Avpr1 or Kiss1impeu neurons (32, 105) | Circadian AVP signaling on Kiss1impeu neurons is facilitated by estrogen during LH surge (32, 105) |
| Corticotropin-releasing hormone (CRH) | Close appositions of CRH-immunoreactive fibers and Kiss1 neurons (129) | 1. Central administration of CRH suppressed LH secretion in rats (117). 2. CRH stimulated LH and increased GnRH pulse amplitude in sheep (118, 119) | CRHNR (125) | Possible regulator of GnRH secretion in stress-induced reproductive disorders |
| Dynorphin (DYN)                 | DYN-immunoreactive fibers in close apposition with Kiss1impeu neurons (mice) (171, 172) | 1. ICV injection of dynorphin in adult male rats suppressed LH secretion (71) | Oprk1 or KOR (89, 99, 73) | Inhibitory |
| Ghrelin                         | NKB-immunoreactive fibers in close apposition with Kiss1impeu neurons (mice) (81) | NKB increased action potential firing of Kiss1impeu neurons via activation of NKR (57–59) | NK1 tachykinin receptor (NK1R), NGK and NGR (58) | Stimulatory |
| Neurokinin B (NKB)              | Pituitary adenylate cyclase-activating peptide (PACAP) | Channel rhodopsin-assisted mapping indicated PACAP neurons from the PMV suppresses onto Kiss1 neurons (mice) (81) | Kiss1impeu neurons relay the stimulatory effects of melanocortin signaling onto the reproductive axis during puberty (134) |
| Relaxin-3                       | Dense relaxin-3-immunoreactive fibers projecting to the ARC (mice) (103) | ICV administration of human relaxin-3 in adult male rats stimulated LH secretion (102) | Rdf1 (82) | Stimulatory |
| RFamide-related peptide-3 (RFPR-3) | RFPR-3-immunoreactive fibers contacted Kiss1 neurons (mice and rats) (93, 90) | 1. ICV administration increased LH secretion in rats (91, 93), whereas it had no effects in ewes. 2. Stimulates LH secretion in male hamsters (94, 95), but inhibits LH release in females (87, 93) | Gpr147 or Nphr1 (53, 90) | Primary central target for the inhibitory action of melanin signal on reproductive function (93) |
| Somatostatin (SST)              | Appositions between SST-immunoreactive fibers and Kiss1 neurons (sheep) (89) | Inhibits episodic LH secretion during anestrus in sheep (84). | Sstr1, 2, 3, and 5 (82) | Inhibitory |
| Substance P (SP)                | SP-immunoreactive fibers in close apposition with Kiss1impeu neurons (monkey) (73) | 1. ICV administration of SP showed increased LH release (87). 2. SP activates Kiss1impeu neurons (59) | Tacr1 or NKR1 (68) | Stimulatory; critical in sustaining reproductive capabilities in female mice (66) |

Italic fonts indicated receptor gene expression identified by in situ hybridization or single cell reverse-transcription PCR. Receptors detected by immunostaining or pharmacological blockade were written in capital letters. ICV, intracerebroventricular; LH, luteinizing hormone; PMV, ventral premamillary nucleus of the hypothalamus; NK1, 2, and 3R, neurokinin-1-3 receptor; Sat1, 2, 3, and 5, somatostatin receptor 1, 2, 3, and 5; Opr147, G protein-coupled receptor 147; Rdp1, relaxin/insulin like family peptide receptor 1; MC4R, melanocortin 4 receptor; Lep, lepin receptor; Adyay1r1, adenylate-cyclase-activating peptide 1 receptor 1; CRHR, corticotropin releasing hormone receptor; Npy1r, Npy2r, and Npy5r, neuropeptide Y receptor 1, 2, and 5; Ghsr or GHSR1A, growth hormone secretagogue receptor type 1a; Nphr1, neuropeptide FF receptor 1; Oprk1 or KOR, opioid receptor, kappa 1; Tacr1, tachykinin receptor 1.
neurons, as this information will improve our understanding of how different cues can be integrated into the HPG axis, complementing the homeostatic regulation of reproductive function and maintaining fertility. Infertility is a global health issue affecting a significant proportion of humanity and is estimated to affect 8–12% of couples worldwide (16, 17). Given that kisspeptin has been successfully used in patients with hypothalamic amenorrhea (18, 19) as well as those with an absence of neurokinin B (NKB) signaling (20), insights into these central pathways will aid in the manipulation of kisspeptin signaling that may be used in the treatment of infertility and reproductive disorders.

**THE ROLE OF KISS1 NEURONS IN THE INTEGRATION OF ENDOCRINE RESPONSES**

In mammals, there are two main populations of neurons in the hypothalamus that synthesize kisspeptin. The first is located in the preoptic area (POA) of sheep and monkeys (21–23), or the rostral periventricular area of the third ventricle (RP3V) in rodents (24, 25), hereafter, termed Kiss1RP3V. Kiss1RP3V neurons show a clear sexual dimorphism with greater numbers present in females (26) where they control the GnRH/LH surge that triggers ovulation (27). The Kiss1RP3V neurons express both estrogen receptor alpha (ERα) and progesterone receptor (27). Activation of Kiss1RP3V neurons by estradiol is essential for the positive feedback action of estrogen on the HPG axis and is associated with increased Kiss1 mRNA expression (28). This is dependent on ERα because the increased firing of Kiss1RP3V neurons in response to estradiol is absent in ERα knockout mice (29–31). Peptides such as arginine vasopressin (AVP) (32) and gonadotropin-inhibitory hormone (GnIH) (33) also regulate Kiss1RP3V neurons. Kiss1RP3V neurons have been shown to co-express dopamine and galanin (34, 35). Neuroanatomical tracing using a recombinant adrenovirus encoding farnesylated enhanced green fluorescent protein (EGFP) to facilitate the labeling of Kiss1 neural axons revealed projection of Kiss1RP3V neurons to GnRH neuronal soma and proximal dendrites within the POA (36). The Kiss1RP3V neurons also project to the arcuate nucleus (ARC) and to the distal dendrites of GnRH neurons, suggesting that Kiss1RP3V and Kiss1ARC communicate with each other to synchronize or coordinate LH secretion (36).

A second population of Kiss1 neurons, which are less well characterized, is found in the ARC region of the brain (also called the infundibular nucleus in humans and primates) (10, 21, 37, 38). In rodents, sheep, and goats, these neurons have been shown to co-express other neuropeptides such as NKB and dynorphin (DYN), and this has led to Kiss1ARC neurons being termed KNDy neurons (39–41). In KNDy neurons, estrogen suppresses Kiss1 expression via ERα (28), and embryonic deletion of Esr1 specifically in Kiss1 neurons, advances puberty onset in association with significantly elevated LH levels (26, 42). Kiss1ARC neurons innervate the distal dendrons (a term describing a single projection structure that functions simultaneously as an axon and dendrite) (43) of GnRH neurons. Changes in Kiss1ARC neuron number, morphology, and connectivity with GnRH neurons have been detected across developmental stages (44–48).

**THE ROLE OF KISS1ARC NEURONS IN NEUROPEPTIDE INTEGRATION**

Optogenetic stimulation of Kiss1ARC neurons in vivo has shown an important role for these neurons in orchestrating pulsatile GnRH/LH secretion (49). Since then, the mechanism by which Kiss1ARC neurons contribute to the GnRH pulse generator has been studied. Navarro and colleagues proposed a model where Kiss1ARC neurons are interconnected and use the co-expressed neuropeptides NKB and DYN (39) to form a synchronized network sending a rhythmic stimulatory signal to the GnRH neurons, thus generating pulsatile gonadotropin secretion. Interestingly, optogenetic inhibition of Kiss1ARC neurons revealed that the middle or caudal Kiss1ARC neurons are responsible for pulsatile LH secretion whereas inhibition of rostral ARC failed to suppress LH release (50). The ARC population has been revealed to have higher number of Kiss1 cells compared to the RP3V region (51, 52), and they may be heterogeneous in terms of firing pattern or ion channel distribution, and possibly receptor expression (53, 54). All in all, we are beginning to appreciate the complexity of the Kiss1ARC neurons in terms of their morphology, functional heterogeneity, different projection areas, and afferent inputs.

Central administration of various classical neurotransmitters and neuropeptides has been found to alter LH secretion through a GnRH neuron-dependent pathway (55). Given that Kiss1ARC neurons are upstream regulator of GnRH neurons, this indicates that Kiss1ARC neurons may possibly receive signals from a variety of neuropeptides and neuromodulators, which modulate GnRH/LH secretion by interacting with receptors located on kisspeptin neuron cell bodies, dendrites, and terminals. Another way to evaluate upstream signals to Kiss1 neurons is to use electrophysiological recordings after neuropeptide stimulation in brain slice preparations and selective pharmacological inhibition of these responses. This approach was taken to show that Kiss1 neurons can respond directly to NKB, DYN (56–58), and substance P (SP) (58). Once an initial firing response is identified, the next step is to undertake intracerebral injection or intracerebroventricular (ICV) infusion of the neuropeptide or antagonist in vivo to assess the corresponding physiological effects.

There is good evidence regarding the effects of the tachykinin NKB on Kiss1ARC neurons and subsequent LH release. Kiss1ARC neurons are depolarized and increase action potential firing upon activation of NK3R, the membrane receptor for NKB (57–59). In rodents, the full excitatory effect of NKB on Kiss1ARC neuron firing requires the activation of all three tachykinin receptor subtypes [NK1 tachykinin receptor (NK1R), NK2R and NK3R], which may all be expressed in these neurons (58). Activation of NK3R, with the agonist senktide, has been used in several studies to probe the effects of NKB on LH release in rodents (39, 60–62). The mechanisms of Kiss1ARC NKB signaling may vary between species as pulsatile LH release is sensitive to NK3R blockade alone in sheep (63), while in rodents, it requires blockade of all three tachykinin receptors with no effect with blockade of NK3R alone (64). The theory of a synchronized KNDy network as the pulse generator may be plausible; however, it is not clear whether these reciprocal KNDy–KNDy connections (65) derive...
from axon collaterals within a single neuron or connections from neighboring KNDy cells, or inputs from a segregated subset of KNDy neurons.

Substance P (SP) encoded by the Tac1 gene, is another tachykinin, which can also influence reproduction. Female Tac1 knockout mice display delayed puberty (66). Early studies with ovarioctomized and estrogen-primed rats treated with intravenous or ICV administration of SP showed increased LH release (67). SP acts via the NKR1, and gonadotropin stimulation is blocked in the absence of kisspeptin (68). Moreover, SP activates Kiss1 neurons (58) and approximately half of the Kiss1 neurons express gene encoding the SP receptor (NKR1) (68). These findings support a role for SP acting via Kiss1 neurons to stimulate GnRH release and its critical role in sustaining reproductive capabilities in female mice. In rodents, populations of substance P cells were found in the ventromedial nucleus of the hypothalamus (VMH) (68, 69). One study reported SP fibers projection to the ARC, and they are in close apposition with Kiss1 neurons in male juvenile monkey (70). Majority of the SP cells were found in the premammillary nucleus, sparse SP cells were also observed in the ARC of the monkey.

Another substance that has been identified as an inhibitor of gonadotropin secretion in DYN (71), which belongs to the family of endogenous opioid peptides and is considered to mediate the negative feedback effects of progesterone on LH secretion (72). It has been reported that DYN exerts its inhibitory effects through Kiss1 neurons (41) and the theory of DYN’s inhibitory effect on the GnRH pulse generator has emerged. More than 90% of Kiss1 neurons in the ewe express kappa opioid receptor (KOR) (73); whereas a lower percentage of KOR was revealed by in situ hybridization (39, 74) and single-cell reverse transcription-PCR studies (59). It is hypothesized that DYN may inhibit GnRH pulse frequency by binding to postsynaptic KOR in Kiss1 neurons. Further elucidation of the mechanism underlying DYN/KOR-dependent GnRH pulse generator suppression is yet to be proven.

The role of pituitary adenylate cyclase-activating peptide (PACAP) in puberty has been shown in several recent studies. In ovarioctomized ewes, ICV administration of PACAP depressed plasma LH amplitude and pulse frequency (75). In contrast, intravenous infusion of PACAP elevated the plasma LH levels in male rats (76). Knockout of the PACAP gene (Adcyap1) is partially lethal (C57Bl/6j genetic background) as the majority of PACAP-deficient mice died at around 3 weeks of age (77) from dysfunction of lipid and carbohydrate metabolism (78). Surviving PACAP-deficient female mice exhibited reduced fertility, with no obvious defects in the length of estrus cycles but their mating frequency was significantly reduced (79). Abundant expression of Adcyap1 is found in the ventral premammillary nucleus of the hypothalamus (PMV) and the VMH, both regions known to be involved in leptin-related control of puberty and fertility (80, 81). Despite the presence of leptin receptor (LepR) in Kiss1 neurons (82), the main site of leptin’s action to regulate reproduction is through cells in the PMV (83). Recently, channel rhodopsin-assisted circuit mapping revealed that the PMV PACAP neurons form direct monosynaptic contact with both Kiss1 and Kiss1 neurons, which express the PACAP receptor (81). Furthermore, calcium-imaging experiments provided intriguing insights that PACAP exerts direct stimulatory effect exclusively on caudal Kiss1 neurons.

The peptide hormone somatostatin (SST), acting through the SSTR2 receptor, inhibits episodic LH secretion possibly via the mediobasal hypothalamus (MBH) during anestrous in sheep (84). Recently, Dufourny and Lomet discovered reciprocal connections between Kiss1 and SST neurons in ewes (85), with most Kiss1 neurons showing SST-immunoreactive fiber appositions. The expression of Sstr2 by Kiss1 neurons is not yet proven but Sstr2, 3, and 4 are expressed in GnRH neurons in mice (86). SST neurons can be found in the periventricular area of the POA, in the ARC and in the ventrolateral area of the VMH. The functional relevance of Kiss1 appositions on SST neurons remains to be verified since GPR54 has not yet been identified in SST neurons.

The RFamide-related peptide-3 (RFRP-3) is a mammalian analog of avian GnIH, found primarily in the dorsomedial nucleus of the hypothalamus (DMH) and adjacent structures (87, 88). This peptide inhibits LH secretion by suppressing GnRH secretion (89). Anatomical studies showed that about 20% of Kiss1 neurons from proestrus female mice were contacted by RFRP-3 fibers, and only a small fraction of the Kiss1 neurons expressed Gpr147, one of the receptors for RFRP-3 (33). Similarly, 35% of Kiss1 neurons receive RFRP fiber contacts, with approximately 25% expressed Gpr147 (90). Even though RFRP-3 is considered an inhibitor of gonadotropin secretion in rats (91, 92) and female hamsters (87, 93), it is able to stimulate the gonadotropic axis in male Syrian and Siberian hamsters. RFRP-3’s stimulatory effects on gonadotropin and testosterone production were observed in male hamsters (94, 95). However, in the ewes, the initial inhibitory effects on LH secretion was contradicted by a study performed by Decourt and colleagues (97), suggesting that RFRP-3 had no direct effect on LH release. Recently, RFRP-3 expressing neurons emerged as the primary central target for the inhibitory action of the melanotin signal after melatonin was reported have little or no effect on Kiss1 neurons (98). RF-amide peptides are highly regulated by the melanotin-driven thyroid-stimulating hormone (TSH), which are critical for the control of seasonal breeding (99).

The neuropeptide oxytocin, which is involved in social bonding and sexual reproduction may also interact with Kiss1 neurons. Central kisspeptin excitation of oxytocin neurons occurs in late pregnancy, and this excitation is likely to be mediated by a subpopulation of Kiss1 neurons projecting to the supraoptic nucleus (100). In contrast, the effect of oxytocin on Kiss1 neurons is unclear. One study revealed that intranasally applied oxytocin reaches the brain and oxytocin treatment at its highest dose increased the level of Kiss1 and NKB mRNA in the anterior hypothalamus of female rats by approximately 400%, but Kiss1 expression was unaffected by the lower doses of oxytocin. Also, elevated Gnrh mRNA expression following intranasally applied oxytocin was observed but the plasma LH concentration remained normal (101). Kiss1 neurons have not been shown to express the oxytocin receptor; hence, the direct impact of oxytocin on Kiss1 neurons within the anterior hypothalamus has yet to be confirmed.

McGowan and colleagues discovered a novel role for the neuropeptide relaxin-3 in the stimulation of the HPG axis via hypothalamic GnRH neurons (102), suggesting that this peptide may
play a role in coordinating feeding and reproductive responses in response to alterations in energy balance. ICV administration of human relaxin-3 (H3) in adult male rats significantly increased plasma LH and the effect was blocked by pretreatment with a peripheral GnRH antagonist. H3 stimulated the release of GnRH from hypothalamic explants and GT1-7 cells, which express relaxin/insulin like family peptide receptor 1 and 3 (RXFP1 and RXFP3). Immunohistochemical labeling of relaxin-3-expressing neurons in male rats indicated dense relaxin-3-immunoreactive fibers projecting to the ARC. These relaxin-3-expressing cells were derived from the nucleus incertus, pontine raphe nucleus, periaqueductal gray, and the dorsal area to the substantia nigra detected by in situ hybridization (103). Further studies are required to define the physiological importance of relaxin-3 in the regulation of reproduction, and the possible interaction with Kiss1<sup>ARC</sup> neurons.

**Kiss1 Neurons and the Integration of Environmental Cues**

Many of the studies investigating Kiss1<sup>RPV</sup> regulation have focused on factors that coordinate the LH surge with environmental cues, such as circadian inputs. Vasopressin (AVP) neurons from the suprachiasmatic nucleus (SCN) are the key SCN neurons regulating Kiss1<sup>RPV</sup> via the V1a receptor and AVP-immunoreactive fibers project to Kiss1<sup>RPV</sup> neurons (32). Vasopressin signaling is critically dependent on estrogen, in the presence of which vasopressin exerts a potent and direct stimulatory influence upon most Kiss1<sup>RPV</sup> neurons (104, 105). Interestingly, estrogen permits circadian AVP signaling at Kiss1<sup>RPV</sup> neurons without changes of AVP receptor signaling throughout the estrus cycle (105).

Seasonal breeders use photoperiod or day length as the primary environmental cue to time reproduction (106), ensuring birth occurs when environmental conditions favor the energetic demands of lactation and survival of offspring. The circulating levels of melatonin synchronize reproduction with photoperiod in these animals (98). These seasonal variations in reproduction are the direct result of changes in the neural network, specifically in (i) upstream neurons sensing the melatonin secretion from the pineal gland in response to photoperiod; (ii) neurons in the hypothalamus regulating GnRH and LH secretion (107). Ewes are short day breeders and remain in anestrous in long day conditions. Conversely, hamsters are long-day breeders (98). Both steroid-dependent and steroid-independent inhibition of gonadotropin secretion corresponding to photoperiod was demonstrated in sheep, Syrian and Siberian hamsters (108, 109). While Kiss1<sup>ARC</sup> neurons are the central site for the negative steroid feedback occurring in the breeding season, Kiss1 expression is also inhibited by the short day melatonin signal (94, 110, 111). In the ewe, long days activate glutamatergic neurons that innervate A15 dopaminergic neurons in the retrochiasmatic area (112). Dopamine released from these neurons inhibits Kiss1<sup>ARC</sup> neuron activity, thus inhibiting GnRH/LH secretion and inducing infertility (113). In Syrian hamsters, long days stimulate the release of RFRP-3-expressing neurons in the DMH (114), which increases the activity of Kiss1<sup>ARC</sup> neurons. It is hypothesized that the increase in kisspeptin then stimulates GnRH secretion, causing testicular recrudescence and the resulting testosterone elevation stimulates Kiss1<sup>RPV</sup> neurons (94). Conversely, kisspeptin appears to play no role in mediating the effects of photoperiod in male Siberian hamsters (115).

The hypothalamic paraventricular nucleus (PVN) contains a prominent population of corticotrophin-releasing factor (CRF) neurons, which regulate the hypothalamic–pituitary–adrenal axis (116). The role of PVN CRF in the control of LH secretion, however, is controversial even though it is known that stress responses can affect fertility. Central administration of CRH suppressed LH secretion in rats (117), whereas this is not the case in sheep (118, 119). Evidence of synaptic connections between CRF and GnRH neurons in the medial POA of rats (120) and the MBH of humans (121) indicate direct functional connection between CRF and GnRH neurons. Nevertheless, tract-tracing studies failed to find any CRF neurons projecting from the PVN to the POA where most GnRH neuronal soma are found in the rat (122). Stress-induced elevated CRF mRNA expression within the PVN did not correlate to LH pulse suppression (123) and finally, PVN lesions failed to interfere with the inhibitory effect of stress on LH release in rats (124). This suggests that CRF may act on one of the regulators of GnRH neurons rather than directly on GnRH neurons. Indeed, double-labeling immunohistochemistry revealed that most Kiss1<sup>RPV</sup> and Kiss1<sup>ARC</sup> neurons in the female rat hypothalamus expressed the CRF receptor, CRHR. Close appositions of CRH-immunoreactive fibers on some of the Kiss1<sup>RPV</sup> and Kiss1<sup>ARC</sup> neurons have been reported (125).

**Metabolic Integration via Kiss1<sup>ARC</sup> Neurons**

Reproduction is normally coordinated with nutritional status to ensure that pregnancy, parturition, and lactation occur during periods of ample food to maximize the survival of the individual and their offspring (126). One way in which this is coordinated is via the anorexigenic and orexigenic actions of proopiomelanocortin (POMC) and agouti-related peptide (AgRP) neurons, respectively within the ARC on Kiss1<sup>ARC</sup> neurons. POMC and AgRP neurons are the first-order sensors of peripheral metabolic signals, such as the leptin and insulin to maintain energy homeostasis. The adipose tissue-derived hormone, leptin and stomach-derived ghrelin have also been implicated to exert effects on Kiss1<sup>ARC</sup> neurons.

**Alpha-Melanocyte-Stimulating (α-MSH) Hormone From POMC Neurons During Puberty**

The role of α-MSH in the control of gonadotropin secretion in adults has been tested pharmacologically (127) and direct effects of α-MSH on GnRH neurons has been shown in adult mice (128). Central alpha-MSH can stimulate or inhibit LH secretion in rats depending on the hormonal milieu (129, 130). Pomc and Kiss1 neurons appear to make mutual contacts in the adult ovine brain, α-MSH enhances Kiss1 mRNA levels in the POA of sheep and decreases Kiss1 expression in the ARC. Kisspeptin has been shown
to inhibit Pomc gene expression in the ARC of the sheep (131), while ARC POMC neurons displayed increased firing after kisspeptin stimulation in mice (132). The cocaine- and amphetamine-regulated transcript (CART), another neuropeptide co-expressed in most POMC neurons can directly stimulate Kiss1ARC neurons in female mice (133), implying a possible functional connectivity between POMC and Kiss1ARC neurons. Manfredi-Lozano and teammates (134) performed a comprehensive pharmacogenetic and optogenetic approaches where using expression analyses, electrophysiological recordings, and a chemogenetic approach to pinpoint the physiological role of leptin acting via an α-MSH-kisspeptin pathway in the metabolic regulation of puberty. Kiss1ABC neurons appear to transmit the stimulatory effects of melanocortin signaling onto the reproductive axis during puberty based on these data: (1) presence of appositions between α-MSH fibers and Kiss1 neuronal cell bodies of pubertal female rats; (2) reduction of Kiss1 mRNA expression in the ARC of pubertal females subjected to chronic inhibition of melanocortin 3 and 4 receptors (MC3/4R); (3) significant attenuation of LH responses to α-MSH in mice with congenital inactivation of Gpr54; and, importantly, (4) reduced LH responses to α-MSH following chemogenetic inhibition of Kiss1ABC neurons.

**Gamma-Aminobutyric Acid (GABA) From AgRP Neurons During Metabolic Deficiency**

A seminal study by Padilla and colleagues indicated that AgRP-expressing neurons are activated during starvation and are involved in leptin-associated infertility during negative energy state (135). Using AgRP-neuron ablation and optogenetic strategies, they discovered inhibitory synaptic connections of AgRP neurons with neighboring Kiss1ARC neurons and rostral Kiss1RP3V neurons. The activated AgRP neurons release GABA, which has direct inhibitory actions on Kiss1ARC neurons. In agreement with this, Kiss1ARC neurons received less pre-synaptic inhibition in the absence of AgRP neurons after neonatal toxin-induced ablation. Chemogenetic activation of AgRP neurons as means of enhancing the activity of AgRP neurons over a sustained period is sufficient to perturb fertility in vivo. As a result, the animals exhibited delayed estrus cycles and decreased fertility (135). Interestingly, a direct, GABA-mediated connection between AgRP and GnRH neurons was not observed in this particular study despite the evidence that GnRH neurons are sensitive to a melanocortin agonist (136) and expressed NPY Y1 and Y5 (NPY1R and NPY5R) receptors (137). These findings confirmed that AgRP signaling contributes to infertility by inhibiting Kiss1 during metabolic deficiency.

**Leptin and Ghrelin**

Leptin deficiency is associated with suppressed Kiss1 expression in rodents and sheep, while leptin administration has been shown to increase Kiss1 expression (131, 138, 139) as well as elevating LH pulse frequency, amplitude, and mean levels (140). Although Kiss1ARC neurons express the LepR, only a small fraction of Kiss1ABC neurons are responsive to leptin (141) and deletion of LepRs from Kiss1 neurons resulted in no puberty or fertility deficits (83). To eliminate the possibility of developmental adaptations and system redundancies, the LepR was selectively re-expressed in Kiss1 neurons of Lepr-null mice. These mice showed no pubertal development and no improvement of the metabolic phenotype: they remained obese, diabetic, and infertile (142). These findings clearly demonstrate that Kiss1 neurons are not the direct target of leptin during puberty onset. Cravo and colleagues also confirmed that leptin signaling in Kiss1 neurons occurs only after completion of sexual development (142).

In addition to its undisputed role in the regulation of metabolism and energy balance, increasing evidence shows that ghrelin can influence fertility. Studies conducted in several species, including rats, sheep, and humans, indicate that ghrelin administration suppresses gonadotropin secretion (143–145). The ghrelin receptor, GHSR1a (growth hormone secretagogue receptor) is present in several hypothalamic regions, including those known to be involved in the control of the reproductive function, indicating that this hormone can interact directly with hypothalamic neurons (146). Work by Frazao and colleagues (147) confirmed that ghrelin interacts directly with a subpopulation of Kiss1ARC neurons to modulate their activity and that exposure to estradiol increases the sensitivity of these neurons to ghrelin signals. The effects of ghrelin varies according to the estrogen milieu, as it exerts a more pronounced orexigenic effect in ovariectomized female rats and diestrus females when estrogen levels are low. Males with estradiol treatment are resistant to the stimulatory effects of ghrelin on food intake (148). The physiological relevance of ghrelin effects on Kiss1ARC neurons in food intake and metabolic regulation requires further investigation.

**EXPERIMENTAL APPROACHES FOR INVESTIGATING FUNCTIONAL INPUTS TO Kiss1ARC NEURONS**

Recently, methodological advances have allowed us to gain significant insights into upstream signals that converge on Kiss1 neurons to modulate the reproductive axis. One approach has been to use of single cell RNA sequencing (scRNA-seq) to identify the gene expression profile of Kiss1 neurons and thereby identify the repertoire of surface receptor that may mediate physiological responses (82, 149). This approach provides the most direct and unbiased method to define a cell type based on its transcriptional profile, which can provide additional insights into connectivity and function (150, 151). Two groups recently carried out single cell analysis of neurons from the hypothalamus or the ARC that included Kiss1ABC neurons (82, 149). Results from these studies provide extensive information about the neuropeptides, neurotransmitters, and receptors co-expressed in Kiss1ABC neurons, facilitating the assessment of crosstalk among different neuropeptide signals within the same cell. Both scRNA-seq studies revealed high expression of Slc17a6, which encodes a vesicular glutamate transporter and Tic2 (NKB) in Kiss1ABC cells. In parallel, with previous data from electrophysiology, in situ hybridization and immunohistochemistry studies, Kiss1ABC neurons have been shown to express the NK3R (Tacr3), estrogen receptor 1 (Esr1), receptors for progesterone, prolactin, ghrelin, and the neuropeptide FF receptor 1 (27, 28, 74, 147, 152,
153). In addition, the nociceptin receptor (Oprl1), melanocortin receptors (Mch3r and Mch4r), NPY receptors (Npy1r, Npy2r and Npy5r), thyrotropin-releasing hormone receptor (Trhr), and insulin receptor substrate 4 (Irs4); are among the receptors that were identified in the Kiss1ARC single cells at low levels (149). These data suggest that Kiss1ARC neurons may receive afferent inputs from neurons involved in nociception, energy homeostasis, insulin signaling, and TSH secretion. Interestingly, given that Kiss1ARC single cells express four SST receptor subtypes (Sstr1, 2, 3, and 5), the PACAP (Adcyap1r1), oxytocin (Oxtr), and RXFP1 (Rxfp 1) receptors (149), this information suggests a possible link between functional relevance of SST, PACAP, oxytocin, relaxin, and reproduction regulation via Kiss1ARC neurons, at least in females.

One limitation of the scRNA-seq studies is that the experiments are often designed to generate gene expression profiles without appropriate considerations of neuroendocrine criteria. Cell samples may be pooled from several animals, combining males and females, and for female samples; the stage of the estrus cycle may not be considered. Nevertheless, these data are still valid as preliminary information about the range of receptors expressed in the Kiss1ARC neural population. Specific scRNA-seq studies are necessary to further characterize the heterogeneity of the Kiss1ARC neurons.

Another approach to investigate functional inputs to Kiss1ARC neurons is to use powerful genetic methods in transgenic mice. The development of transgenic mouse lines with deletion of specific receptors in Kiss1 neurons is a great tool in addressing the physiological relevance of these receptors. For example, mice have been generated in which the Esr1 gene has been ablated in Kiss1 neurons via a CRE-mediated recombination event (51), and these mice have defined the importance of sex steroid signaling in Kiss1 neurons. Acute ablation of a gene from the earliest developmental time point, however, can sometimes be associated with compensatory changes in gene expression that can mask the effect of the gene disruption (154). If this occurs, a better alternative is to create a mouse model with inducible gene disruption to delineate the time windows in which gene inactivation is critical for the functional manifestation of a particular effect (155). Furthermore, varying the onset of gene manipulation at different time points and combining genetic manipulation with pharmacological or behavioral interventions will help to clarify gene–environment interactions that are crucial for the development or maintenance of reproductive phenotypes. The CRISPR/Cas9-based genome-editing tool implemented in mammalian cells has revolutionized gene-editing techniques (156). While this technique has generated huge impact on in vitro studies, progress is being made to also apply it in vivo. The combination of adeno-associated virus (AAV) and CRISPR/Cas9 system may be particularly useful in the future for editing reproductive-associated genes given that encouraging results have been obtained with the next-generation synthetic AAV capsids in several transgenic mouse models (157, 158).

A main objective in deciphering the neural circuitry is to define the synaptic inputs and outputs of specific neuronal sub-populations in different regions. Mapping the network of Kiss1 neuronal inputs and outputs using a combination of molecular genetics and viral tract tracing techniques provides both anatomical and functional circuit information. Until recently, the input–output relationships have been mapped using neuroanatomical tracers to reveal connections between regions (159). Classical tracers, such as biotin-dextran amine, fluorescent latex microspheres, fluorescent cholera toxin conjugates, or phaseolus vulgaris-leucoagglutinin, have provided very useful information to trace fibers in anterograde and retrograde directions, depending on the type of tracer applied (65, 160, 161), but they reveal only the axonal projections, not synaptic connections, and can be difficult to genetically target to specific neuronal types. Trans-synaptic tracing using retrograde viruses such as pseudorabies virus (PRV) is useful in revealing the brain regions forming synaptic connectivity with Kiss1ARC neurons (162), but this technique is only limited to rodents. The PRV is contagious to domestic mammals as it causes Aujeszky disease in cattle and swines. However, the major drawback of this technique is that the PRV crosses multiple synapses, making it difficult to distinguish the first order and higher order synaptic inputs unless the PRV is combined with other neuronal tracers. To overcome the limitations of PRV tracing, monosynaptic tracing using glycoprotein (G)-deleted rabies virus is now a widely adopted method to delineate brain-wide monosynaptic connectivity (163).

Intensive efforts are being made to delineate the complete wiring diagram or connectome of the mammalian brain as a means to better understand how neural circuits control behavior. High-throughput electron microscopy has been used to delineate microscale connectivity (164), while tracing strategies utilizing viral tracers encoded with fluorophores have allowed for milliscal circuit mapping (165). These studies have elegantly dissected a number of complex circuits. However, these methodologies are not designed to provide molecular information about the pre-synaptic neural populations. These shortcomings warrant the identification of marker genes for neurons within the circuits to enable the testing of their functional role.

While neuroanatomical methods enable high-resolution mapping of neural circuitry, these approaches do not allow molecular profiling of neurons based on their connectivity. An advanced approach for translational profiling of neurons based on connectivity using viral translating ribosome affinity purification (vTRAP) has been reported recently (166). In this approach, CRE-dependent AAV or other retrograde viruses (rabies or canine adenovirus) are engineered to express an EGFP-tagged ribosome protein enabling isolation of mRNA that is being translated from a discrete CRE-expressing neural population. Projection-specific translational profiling is achieved by selectively precipitating neuronal ribosomes based on connectivity. Quantitative PCR is then used for selected target genes or high-throughput RNA sequencing to determine the neuronal identity without the need for detailed anatomical or electrophysiological investigation (166). The drawback of this technique is that high-throughput RNA sequencing on the immunoprecipitated RNA is critical and a substantial amount of validations is required prior to selection of appropriate marker genes for the projecting neurons.

An elegant study performed by Nectow and colleagues also used the vTRAP method to delineate the dorsal raphe nucleus (DRN) circuit in regulating feeding (167). First, they used an unbiased approach to map sites of neural activation in response...
to fasting, re-feeding, and hormonal cues, where subsets of DRN neurons were activated. Following this, comprehensive pharmacogenetic and optogenetic approaches were applied to carefully dissect the roles of DRN GABAergic neurons (DRN\textsuperscript{VGLUT3}) and glutamatergic neurons (DRN\textsuperscript{VGLUT1}) in modulating food intake. Transmembrane receptors that were enriched in both DRN\textsuperscript{VPA} and DRN\textsuperscript{VGLUT3} neurons using the vTRAP method can be used for pharmacological screening of the ligands of these receptors. Finally, the effects of appropriate ligands for each receptor associated with the predicted response on feeding were tested using electrophysiological studies. This approach could also be used to molecularly characterize how the transcriptional profile of Kiss1\textsuperscript{ARC} neurons changes in response to physiological stimuli.

Another powerful method to identify functional inputs into Kiss1 neurons is to use optogenetic or chemogenetic approaches. These involve a priori identification of the specific pre-synaptic neural population as these methods required the use of neuron-specific CRE-expressing animal model. Both approaches involve the expression of non-native proteins that can function as channels (channel rhodopsin), pumps, or receptors in neurons. These novel proteins are sensitive only to exogenous non-native stimuli such as light or the compound clozapine-N-oxide (CNO). In this way, both approaches provide an exclusive selectivity for neuronal manipulations, enabling causal analyses of the roles of neural circuits in defined functions (168). Optogenetics allows millisecond-scale temporal accuracy in manipulating neuronal activity, which is critical for assessing circuit or behavioral functions that emphasizes on the rate or timing of neural activity. However, light activation of channel rhodopsin expressing neurons mainly release fast neurotransmitters such as glutamate and GABA; whereas neuropeptides require higher frequency and prolonged stimulations to be released. These factors may have explained the apparent lack of success for optogenetic release of neuropeptides (169). Alternatively, designer receptors exclusively activated by designer drugs that are sensitive to CNO and can be stimulated by a simple systemic CNO administration can be used. The main limitation of this approach is that their action is slow, in the order of minutes, rendering limited application in analyzing neural processes that rely on rate or timing of neural activity. But, this is ideal for examining the effects of chronic stimulation of neuronal populations.

To conclude, the plasticity or dynamics of the underlying kisspeptin–GnRH network in different physiological conditions is important. Relatively little is known about the roles of kisspeptin under various physiological conditions such as negative energy state, lactation, and reproductive senescence. An excellent way to investigate the dynamics of the neural circuits connecting to Kiss1\textsuperscript{ARC} neurons under these physiological conditions is to implement vTRAP on Kiss1\textsuperscript{ARC} neurons followed by extensive chemogenetic and optogenetic strategies for causal analyses of the roles of specific neural circuits in defined behavioral responses or reproductive phenotypes.

**CONCLUDING REMARKS**

Precise control of gonadotropin release by the HPG axis is essential for sustaining fertility in all mammals (170). Therefore, the HPG axis must be able to respond to changes in endocrine, metabolic, or environmental cues to regulate GnRH/LH release. The Kiss1\textsuperscript{ARC} neurons are positioned as an ideal hub receiving afferent inputs from other brain regions in response to the internal homeostatic and external signal. Our understanding of the neural networks connecting with Kiss1\textsuperscript{ARC} neurons is limited, however. To precisely identify and functionally characterize specific synaptic inputs to Kiss1\textsuperscript{ARC} neurons poses a challenge, given that: (1) Kiss1\textsuperscript{ARC} neurons are heterogeneous; (2) estrogen may have an organizational effect on the inputs; (3) there could be an interplay between the Kiss1\textsuperscript{ER\textsuperscript{3}} and Kiss1\textsuperscript{ARC} neurons in fine-tuning pulsatile GnRH/LH release. The search for afferent inputs into Kiss1\textsuperscript{ARC} neurons is continuing using new technologies to decipher the neural network associated with the Kiss1-GnRH system. Techniques such as trans-synaptic viral tracing, single cell RNA sequencing combined with optogenetics and chemogenetics to allow functional analyses are providing significant knowledge about the regulation of Kiss1\textsuperscript{ARC} neurons. The precise mechanisms delineating how neuropeptides/neuromodulators regulate Kiss1\textsuperscript{ARC} neurons and fine-tune GnRH/LH secretion requires further characterization and validation. Also, careful considerations need to be implemented to distinguish direct actions of the neuropeptides on GnRH neurons or effects mediated through Kiss1\textsuperscript{ARC} neurons.

**AUTHOR CONTRIBUTIONS**

S-HY wrote the review with input from WC.

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