Investigating the NPY/AgRP/GABA to GnRH neuron circuit in prenatally androgenized PCOS-like mice

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Abstract

Polycystic ovary syndrome (PCOS), the most common form of anovulatory infertility, is associated with altered signalling within the hormone-sensitive neuronal network that regulates gonadotropin-releasing hormone (GnRH) neurons, leading to a pathological increase in GnRH secretion. Circuit remodelling is evident between GABAergic neurons in the arcuate nucleus (ARN) and GnRH neurons in a murine model of PCOS. One third of ARN GABA neurons co-express neuropeptide Y (NPY), which has a known yet complex role in regulating GnRH neurons and reproductive function. Here, we investigated whether the NPY-expressing subpopulation (NPY\textsuperscript{ARN}) of ARN GABA neurons (GABA\textsuperscript{ARN}) is also affected in prenatally androgenized (PNA) PCOS-like NPY\textsuperscript{ARN} reporter mice [Agouti-related protein (AgRP)-Cre;τGFP]. PCOS-like mice and controls were generated by exposure to dihydrotestosterone or vehicle (VEH) in late gestation. τGFP-expressing NPY\textsuperscript{ARN} neuron fiber appositions with GnRH neurons and gonadal steroid hormone receptor expression in τGFP-expressing NPY\textsuperscript{ARN} neurons were assessed using confocal microscopy. Although GnRH neurons received abundant close contacts from τGFP-expressing NPY\textsuperscript{ARN} neuron fibers, the number and density of putative inputs was not affected by prenatal androgen excess. NPY\textsuperscript{ARN} neurons did not co-express progesterone receptor or estrogen receptor α in either PNA or VEH mice. However, the proportion of NPY\textsuperscript{ARN} neurons co-expressing the androgen receptor was significantly elevated in PNA mice. Therefore, NPY\textsuperscript{ARN} neurons are not remodelled like the wider GABA\textsuperscript{ARN} population by prenatal androgen excess, indicating GABA-to-GnRH neuron circuit remodelling occurs in a presently unidentified non-NPY/AgRP population of GABA\textsuperscript{ARN} neurons. NPY\textsuperscript{ARN} neurons do, however, show independent changes in the form of elevated androgen sensitivity.

Keywords: Neuropeptide Y; steroid hormone receptors; transgenic mouse; tract-tracing; HPG axis
Introduction

Polycystic ovary syndrome (PCOS) is the most common form of anovulatory infertility, affecting 8-10% of women of reproductive age (1). While it is typically described as a highly heterogeneous disorder, neuroendocrine impairment is a consistent feature among women with PCOS. Between 50-75% of women with PCOS exhibit evidence of luteinizing hormone (LH) hypersecretion (2-4), and 90% have an elevated ratio of LH to follicle-stimulating hormone secretion (4). Serial blood sampling shows that this hypersecretion reflects a persistently elevated LH pulse frequency (5,6), indicating a net increase in the activity of the gonadotropin-releasing hormone (GnRH) neural network. This elevated hypothalamic output is likely to stem in part from diminished negative feedback by ovarian steroid hormones, as exogenous estradiol and progesterone are less effective at reducing LH secretion in women with PCOS (7,8). Androgen excess, a key feature of PCOS, may impede negative feedback, as long-term blockade of the androgen receptor in women with PCOS can restore sensitivity to ovarian steroid hormones (9). Such observations have spurred research in animal models using androgen excess to recapitulate PCOS-like features (10) to determine the loci of disrupted hormone sensitivity and associated circuit alterations within the GnRH neural network.

Functional and anatomical findings in both ovine and murine models of PCOS suggest that altered afferent GABAergic input to GnRH neurons may play a role in elevated GnRH/LH secretion (11-19). Prenatally androgenized (PNA) mice that reflect the cardinal features and neuroendocrine impairments of PCOS (20), display a greater frequency of GABAergic post-synaptic currents in GnRH neurons (13,15) and elevated GnRH neuron firing frequency (15,19,21) compared with fertile controls. This is associated with a greater number of closely associated presynaptic GABAergic terminals and a dramatically increased projection of GABAergic fibers from the arcuate nucleus (ARN) (14). Elevated GABAergic input to GnRH neurons appears to be due to early network organisation, as both pre-synaptic markers for GABAergic terminals and post-synaptic GABA currents are elevated by three weeks of age (16,22). The ARN GABA neuron population (GABA_{ARN}) in PNA mice also exhibits a reduction in progesterone receptor expression, indicating reduced sensitivity to this important feedback cue (14). Specific activation of the GABA_{ARN}-to-GnRH neuron circuit with optogenetic and chemogenetic tools elicits LH secretion and mimics some features of PCOS, such as disrupted ovulatory cycles, reduced presence of corpora lutea in the ovary,
and in increase in circulating testosterone (17), suggesting that modifications in this circuit may underpin the elevated LH secretion evident in PNA-treated, PCOS-like mice.

GABA<sub>ARN</sub> neurons in rodents are a large heterogeneous population of neurons that co-secrete a range of neuropeptides and transmitters with implicated roles in the control of GnRH neurons (23-31). What has remained unclear is whether PNA-induced circuit alternations occur in GABA<sub>ARN</sub> neurons as a whole, or within particular subsets of the population. Neuropeptide Y (NPY) is co-expressed in one third of the GABA<sub>ARN</sub> population in both fertile and PNA-treated female mice, and nearly all NPY neurons in the ARN are GABAergic (31). NPY, along with co-expressed agouti-related protein (AgRP), is a well-established regulator of energy homeostasis (32), that is also highly implicated in the regulation of the reproductive axis (33-36). NPY can exert direct effects upon GnRH neurons (37,38), while selective activation of ARN NPY/AgRP neurons regulates upstream kisspeptin neurons via both GABA (39) and NPY receptor-dependent mechanisms (36). Additionally, selective activation of ARN NPY (NPY<sub>ARN</sub>) neurons can cause potent modulation of GnRH/LH secretion (35). Anatomical evidence suggests that NPY<sub>ARN</sub> neurons innervate the proximal region of GnRH neurons (40) and that synapses formed by NPY<sub>ARN</sub> neurons in these proximal regions are GABAergic in nature (40,41). Given that elevated GABA<sub>ARN</sub> input is largely to the proximal dendrite of GnRH neurons in PCOS-like animals (12,14) and the high degree of NPY co-expression in GABA<sub>ARN</sub> neurons (31), we aimed to investigate whether the NPY/AgRP-specific subpopulation of GABA<sub>ARN</sub> neurons is remodelled in the PNA mouse model of PCOS. This was achieved using transgenic AgRP-Cre;τGFP reporter mice to specifically visualise NPY<sub>ARN</sub> cell bodies and fiber projections. We hypothesised that the NPY<sub>ARN</sub>-to-GnRH neuron circuit and the steroid hormone sensitivity of NPY<sub>ARN</sub> neurons would be impacted in PNA-induced PCOS-like mice.
Methods

Animals and tissue collection

The following procedures were carried out with permission from the University of Otago Animal Ethics Committee (Dunedin, New Zealand). Adult female mice were generated and housed in the University of Otago Biomedical Research Facility. Mice were kept in individually ventilated cages, in a climate-controlled environment (20°C/40% humidity) on a 12:12 hour light:dark cycle. Mice were provided ad libitum access to food and water.

Mice in which ARN NPY neurons were identified by green fluorescent protein (GFP) were generated by crossing agouti-related peptide (AgRP)-IRES-Cre mice (42) with ROSA26-CAGS-τGFP floxed-stop reporter mice (43) to generate AgRP-Cre;τGFP mice. Prenatal androgen excess treatment was performed as described previously (13,20), by injection of dams on days 16, 17 and 18 of pregnancy with 100 µL of sesame oil alone (vehicle controls, VEH) or containing 250 µg of dihydrotestosterone (DHT) (prenatally androgenized, PNA). Induction of a PCOS-like phenotype as assessed by daily vaginal cytology smears for 14-days to ensure PNA-treated mice were acyclic (Supplemental Figure 1). Female offspring were studied in adulthood (60-90 days) during diestrus, assessed by vaginal cytology. Following a lethal dose of pentobarbital (3 mg/mL, 100 µL intraperitoneal), animals underwent transcardial perfusion with 4% paraformaldehyde to fix the brain. The brain was then dissected from the skull, cryoprotected in 30% sucrose and cut on a freezing microtome into 30-µm thick coronal sections.

Experiment 1: Assessing NPY\textsuperscript{ARN} neuron projections to GnRH neurons

Double-label fluorescent immunohistochemistry: To assess NPY\textsuperscript{ARN}-to-GnRH neuron circuitry, free-floating immunohistochemistry was performed on every third coronal section through the rostral forebrain including the medial septum (MS; bregma +1.34 to +0.74 mm), rostral preoptic area (rPOA; bregma +0.74 to +0.38 mm) and anterior hypothalamic area (AHA; bregma +0.38 to -0.46 mm) populations of GnRH neurons from VEH (n=5) and PNA (n=8) mice. GnRH neurons were labelled using a guinea pig anti-GnRH primary antibody (1:5,000; GA2, kindly gifted by Prof Greg Anderson; RRID:AB_2721118) (44) and NPY\textsuperscript{ARN} neuron fibers were labelled for the τGFP reporter using a chicken anti-GFP primary antibody (1:5,000; Aves Labs, OR, USA; RRID:AB_10000240) (45). GFP-positive fibers were amplified using a goat anti-chicken AlexaFluor488 antibody (1:200; Molecular...
Prob, OR, USA), while GnRH neurons were visualised using a goat anti-guinea pig AlexaFluor568 antibody (1:200; Molecular Probes, OR, USA). Specificity of secondary antibodies was assessed by primary antibody omission in negative control tissue sets. Mounted sections were coverslipped with Fluoromount G (ThermoFisher Scientific, MA, USA) and kept in the dark at 4°C until imaging.

**Image acquisition and analysis:** Confocal microscopy was performed using a Nikon A1R multi-photon microscope (Nikon Instruments Inc., Melville, NY, USA) to collect images of individual GnRH neurons in the MS, rPOA and AHA. As reported previously (14,16,18), in each animal, z-stack confocal images were collected from five GnRH neurons in the MS and AHA, and ten GnRH neurons in the rPOA, reflecting their distribution density. Using a Plan-Neofluar 40X oil objective (1.30 NA) and 3X digital zoom, scans throughout the soma and the first 75 µm of the primary dendrite of each GnRH neuron were performed using 0.5 µm z-intervals. Pinhole size was maintained at 1 AU using a consistent laser power across animals, while digital gain and offset of red and green channels was kept consistent (<5% variation) to prevent imaging artefacts from confounding or biasing later analysis.

Images of individual GnRH neurons and surrounding GFP-expressing NPY^{ARN} neuron fibers were analysed using NIS Elements software (Nikon Instruments Inc.). The soma circumference in each neuron was measured using pre-established pixel-to-µm conversion preprogrammed into the software; each soma was measured using the image in the stack where the soma was at its largest, and this measurement was recorded. Each primary dendrite was divided into 15 µm segments up to 75 µm, using the measurement tool and manual demarcation of each segment. The number of GFP-expressing fiber contacts associated with each GnRH-neuron was recorded, along with the location of contact (soma or individual dendrite segments). A contact was defined as the point where the red GnRH and green GFP label were contiguous without intervening black pixels, and required that this was present in both the XY plane of view and the orthogonal YZ view of a single focal plane. When a fiber passed across the soma or a segment of dendrite, or ‘bundled’ with the neuron before projecting away, this was recorded as one point of contact. These data were used to calculate the total number of contacts at the level of the soma, primary dendrite as a whole, and within 15µm segments of the dendrite. To calculate the density of appositions, the total number of contacts at the soma were divided by the soma circumference, and the number of contacts within each segment of dendrite were divided by 15, giving a density in
contacts/µm. This provided a measurement of the putative innervation density to each GnRH neuron.

Experiment 2: Assessing steroid hormone receptor expression in NPY\textsuperscript{ARN} neurons

\textit{Double-label fluorescent immunohistochemistry:} To investigate co-expression with gonadal steroid hormone receptors, free-floating immunohistochemistry was performed on coronal sections throughout the rostral, middle and caudal ARN (rARN, mARN, cARN). NPY\textsuperscript{ARN} neuron somata were labelled for GFP reporter expression using a chicken anti-GFP primary antibody (1:5000; Aves Labs) (45) as above. Every third section was co-labelled for one of the following steroid hormone receptors: progesterone receptor (rabbit anti-PR, 1:100; Dako (Agilent), CA, USA; VEH n=8, PNA n=10; RRID:AB_2315192) (46), estrogen receptor \(\alpha\) (rabbit anti-ER\(\alpha\), 1:5000; Millipore; VEH n=5, PNA n=5, RRID:AB_310305) (47), or androgen receptor (rabbit anti-AR PG-21, 1:200; Millipore, MA, USA; VEH n=4, PNA n=4; RRID:AB_310214) (48). To amplify GFP signal, a goat anti-chicken AlexaFluor488 antibody was used (1:200; Molecular Probes, OR, USA), while steroid hormone receptors were labelled using a goat anti-rabbit AlexaFluor568 antibody (1:200; Molecular Probes, OR, USA).

\textit{Image acquisition and analysis:} Confocal microscopy was performed using a Zeiss LSM710 upright microscope (Carl Zeiss AG, Oberkochen, Germany). Confocal z-stacks of two representative sections of the rARN, mARN and cARN were collected from each animal in each experiment using a PlanApo 20X air objective (0.80 NA) to capture one hemisphere in the visual field, using a 2.12 µm z-interval with a pinhole size of 1 AU. High power images for illustrative purposes were collected using a PlanApo 40X oil objective (1.30 NA) with 2x digital zoom to resolve individual soma, using a 0.5 µm z-interval with a pinhole size of 1 AU.

Images were analysed using ImageJ software (National Institutes of Health, Bethesda, MD, USA). The number of cell bodies expressing GFP in each 20X image within each respective stack (z-depth 15-25 µm) was counted, along with the number of cells positive for either PR, ER\(\alpha\) or AR, and finally the number of double-labelled cells in the visual field (a unilateral hemisphere of the ARN). Using this, the percentage of GFP-positive ARN NPY neurons co-labelled with each receptor was calculated.
Statistical analysis

Statistical analysis was performed using PRISM software (Graphpad Software Inc., LA Jolla, CA, USA). Normality of data was assessed by Shapiro-Wilk tests prior to statistical comparisons between VEH- and PNA-treatment groups. Where values from the entire ARN were grouped, VEH and PNA group means were compared using two-tailed unpaired Student’s t-tests to compare absolute values, or Mann-Whitney U tests to compare percentage means. Where VEH and PNA group means were compared in the rARN, mARN and cARN separately, a two-way ANOVA was used, with post-hoc analysis performed using Bonferroni’s multiple comparisons tests. Mean number and density of appositions onto GnRH neurons was compared between VEH and PNA groups using a one-way ANOVA, and post-hoc analysis was performed using Tukey’s multiple comparisons tests. P-values < 0.05 were considered statistically significant.

Results

**NPY**<sub>ARN</sub> projections to GnRH neurons are unaffected in PNA mice

Vaginal cytology, collected daily for 2 weeks prior to tissue collection, demonstrated that all PNA mice were acyclic, spending the majority of time in persistent diestrus and never exhibiting a proestrus smear [Supplemental Figure 1 (49)] as expected. In contrast, all VEH control mice cycled normally, completing 1.6 ± 0.2 full estrous cycles (quantified as proestrus day to proestrus day), and spending 18.8 ± 1.3% of the time in proestrus.

Dense collections of GFP immunoreactive (-ir) NPY<sub>ARN</sub> neuron fibers were observed in close proximity to GnRH neurons located in the rPOA and AHA of both groups (Figure 1A,ii-iii, B,ii-iii). In contrast, NPY<sub>ARN</sub> neuron fibers were not as abundant in the MS of either group (Figure 1A,i, B,i). Of the GnRH neurons imaged across both groups, 117/120 in the rPOA and 56/60 neurons in the AHA received at least one close apposition from a GFP-ir fiber, while just 3/60 neurons in the MS received any close appositions.

The mean number of close appositions per GnRH neuron in each region, compared by two-way ANOVA, was not different between VEH- and PNA-treated groups (F(1, 66) = 0.32, p = 0.57; Figure 1C). Likewise, the mean density of contacts to GnRH neuron somata and primary dendrites in the rPOA, compared using a one-way a ANOVA, was not different between between VEH and PNA mice (F(1, 66) = 0.027, p = 0.87; Figure 1D).
Assessing steroid hormone receptor expression in NPY<sub>ARN</sub> neurons in PNA mice

Androgen receptor: AR-ir was evident in the nuclei and cytoplasm of neurons scattered throughout the ARN of both VEH- and PNA-treated female mice (Figure 2A, 2B), including within the ventromedial subregions where NPY neurons reside. High magnification images revealed AR-ir co-localised within NPY neurons, where it was observed as low-level labelling in the nucleus as well as bright puncta aggregated within the cytoplasm (Figure 2A, 2B). The number of GFP-expressing NPY neurons was not different between in the ARN of VEH- and PNA-treated mice (VEH 229.3± 8.18 cells vs. PNA 246.4 ± 30.04 cells; Figure 2C, Table 1). Although total numbers of AR-ir cells in the whole ARN were not significantly different between VEH- and PNA-treated mice (VEH 598.4 ± 44.79 cells vs. PNA 724.4 ± 32.35 cells; p = 0.067; Figure 2D), a greater proportion of NPY<sub>ARN</sub> neurons were identified to co-express AR in PNA-treated mice (33.2 ± 5.3%) compared with VEH-treated mice (18.9 ± 1.8%, p = 0.045; Figure 2E). No significant differences in AR-ir co-expression were identified in specific ARN subregions (Table 1), however, the number of AR-ir cells in the rostral subdivision of the ARN was significantly elevated in PNA mice (Table 1).

Progesterone receptor: Nuclear PR-ir was found predominantly in the dorsomedial and ventrolateral subregions of the ARN, with very few stained nuclei evident in the ventromedial regions where NPY neuron somata are present (Figure 3A). PR-ir was less abundant and less intense in PNA-treated mice (Figure 3B) compared with VEH-treated mice (Figure 3A). The mean total number of GFP-ir NPY neurons was not different between VEH-treated and PNA-treated mice in the ARN as a whole (Figure 3C), nor in any specific rostral to caudal zone (Table 1). Significantly fewer PR-ir cells were evident in PNA-treated mice through the ARN (336.2 ± 17.11 cells) compared with VEH-treated mice (414.5 ± 6.80; p = 0.0013; Figure 3D). When the rARN, mARN and cARN were compared separately, two-way ANOVA indicated an effect of treatment on the number of PR-ir cells (F(1, 48) = 13.07, p = 0.0007; Table 1), and post-hoc analysis indicated significantly fewer PR-ir cells specifically in the rARN of PNA mice (Table 1). The co-expression of PR within NPY<sub>ARN</sub> neurons was almost entirely absent [0.38 ± 0.2% in VEH-treated mice, and 0.25 ± 0.1% of NPY neurons in PNA-treated mice co-expressing PR (Table 1)].
Estrogen receptor alpha: Images collected throughout the ARN displayed typical ventromedial localisation of GFP-ir NPY neurons, whereas nuclear ERα-ir was scattered around the entire ARN (Figure 4A, top). High magnification images (Figure 4A, bottom) revealed that while ERα-ir nuclei lay in close proximity to NPY^ARN neuron cell bodies, they did not co-localise with NPY^ARN neurons. The mean total number of GFP-ir NPY neurons was not different between VEH-treated and PNA-treated mice in the ARN as a whole (Figure 4B), nor in any specific rostral to caudal zone (Table 1). No difference in the expression of ERα was present between VEH- and PNA-treated mice (VEH 530.9 ± 15.99 cells vs. PNA 559.7 ± 11.15 cells; Figure 4C) within the ARN, nor in any particular rostral to caudal region (Table 1). The co-expression of ERα within NPY^ARN neurons was entirely absent in VEH-treated mice, and extremely limited in PNA-treated mice (0.15 ± 0.08%, Table 1).

Discussion

The present study assessed the impact of prenatal androgen excess, that models PCOS features, on the NPY^ARN-to-GnRH neuron circuit. Using transgenic reporter expression specific to NPY^ARN neurons, we identified extensive NPY^ARN neuron projections to GnRH neurons, particularly to those in the rPOA and AHA. Confocal analysis of close appositions between NPY^ARN neurons fibers and GnRH neurons found no differences in the NPY^ARN-to-GnRH neuron projection in PCOS-like PNA females. These findings suggest that this subset of GABA^ARN neurons are distinct to those that are remodelled by PNA (14,16,18). Although NPY^ARN neurons were found to have virtually no co-expression with PR and ERα, irrespective of prenatal treatment, we did observe a greater proportion of NPY^ARN neurons co-expressing AR in PNA-treated mice, suggesting an upregulation of AR in NPY^ARN neurons in the PCOS-like condition. These finding indicate that although the NPY^ARN population are sensitive to androgens in adulthood, the NPY^ARN-to-GnRH anatomical circuit is not obviously remodelled following prenatal androgen excess.

To dissect NPY/AgRP neurons and their full projections specifically originating from the ARN, AgRP-Cre mice were crossed with a line promoting Cre-dependant τGFP expression (42,43). This was an attractive approach as AgRP and NPY are highly co-expressed in the ARN, and AgRP is exclusively expressed here (50-52). This transgenic model is characterised as both highly specific and highly effective for identifying NPY^ARN
neurons and their projections (35). In addition, this study utilized a well-characterized model of PCOS suited to the study of the neuroendocrine pathology of PCOS (10). This model exhibits the core reproductive abnormalities of PCOS, such as hyperandrogenaemia and anovulation, diminished ovarian hormone negative feedback, and LH hypersecretion (14,20), without the associated metabolic syndrome (53,54). This allows for the characterisation of circuit alterations that result from programmed androgen excess associated with reproductive function, without the confounding co-morbid factors associated with obesity and hyperinsulinaemia present in other models of PCOS (10). However, given the important role of NPY in energy balance, it would be of interest to investigate this circuit in models exhibiting the metabolic phenotype of PCOS.

The vast majority of GnRH neurons within the rPOA and AHA subpopulations received close contacts from NPY\textsuperscript{ARN} fibers, while only 5% of GnRH neurons in the MS subpopulation received NPY\textsuperscript{ARN} fiber contacts. In contrast, tract-tracing of the whole GABA\textsuperscript{ARN} population found approximately half of the MS GnRH neurons receiving close appositions (14). This suggests that GABA\textsuperscript{ARN} subpopulations have different projection patterns with respect to the specific GnRH neurons they innervate. This differential pattern of innervation to the GnRH neurons does not appear to have been previously reported, and supports an ongoing yet unproven hypothesis that GnRH neurons at different regions along their anatomical axis represent functional subpopulations. For example, c-Fos expression suggests that the activity and plasticity associated with surge generation is restricted to the rPOA subpopulation (55,56) and receptor expression studies support the notion that the most rostral MS population may be differentially regulated for distinct (57,58).

Putative innervation of GnRH neurons by NPY\textsuperscript{ARN} neurons was strikingly similar between PNA and VEH groups regardless of GnRH neural subpopulation or whether the soma or dendrite was examined. This analysis was refined further, where the density of contacts was assessed to normalise for variations in soma circumference between GnRH neurons, and by looking at 15 µm sections of the primary dendrite. In this instance, again, there were no differences between VEH and PNA groups. The proximal GnRH neuron dendrite was the focus of this study as GABAergic remodelling is restricted to this region (14,16). However, we cannot rule out changes at the more distal GnRH neuron dendron, the region known to be critical in driving pulsatile LH secretion (59). In any case, the present finding stands in contrast to the plasticity that has been observed in the GABA\textsuperscript{ARN}-to-GnRH neuron circuit, despite NPY\textsuperscript{ARN} neurons composing a large proportion of this circuit (14,31).
This demonstrates that another, as yet undefined population of GABA\textsuperscript{ARN} neurons must be undergoing plastic reorganisation as a result of PNA-treatment, while NPY\textsuperscript{ARN} neurons are a distinct, unchanged population.

Increased GnRH neuronal activity in PCOS-like PNA mice (15,19,21) may reflect a modified balance in the excitatory and inhibitory afferent input that GnRH neurons receive. While GABAergic signalling to GnRH neurons is largely excitatory through GABA\textsubscript{A} receptors (60), and selective optogenetic and chemogenetic activation of GABA\textsuperscript{ARN} inputs to GnRH neurons promotes LH secretion (17), NPY and AgRP likely promote inhibition of the GnRH neurons and LH release. NPY has been shown to act via the Y\textsubscript{1} or Y\textsubscript{4} receptors to inhibit or excite GnRH neural firing rate (38), however NPY binds to Y\textsubscript{1}R at a far greater affinity than Y\textsubscript{4}R (61,62), so the likely net effect is inhibitory. At the cellular level, AgRP has been shown to be stimulatory to a small population of GnRH neurons (38), but also to block the excitatory effects of melanocortin receptor agonists (63). In ovariectomized monkeys, AgRP administration suppresses LH pulsatility (64). In addition, optogenetic and chemogenetic activation of NPY\textsuperscript{ARN} neurons inhibits LH secretion and slows LH pulse frequency in castrate animals (35). Therefore, the absence of enhanced input from the NPY\textsuperscript{ARN} subpopulation of GABA\textsuperscript{ARN} neurons to GnRH neurons in PNA mice aligns with their hyperactive HPG axis state.

Androgens can modulate the development of the NPY\textsuperscript{ARN} neuron population. Male mice and rats have a larger population of NPY\textsuperscript{ARN} neurons compared to females (31,65). Likewise, female ewes treated with T or DHT exhibit greater numbers of AgRP (thus, presumably NPY) neurons in the mARN (66), and prenatal androgen treatment in female ewes also produces greater AgRP fiber density in the POA and other hypothalamic areas (66). NPY/AgRP neurons are implicated in neurodevelopmental processes. However, as shown here and reported previously (31), prenatally androgenised mice do not show any differences in NPY/AgRP cell numbers. The absence of NPY\textsuperscript{ARN} remodelling in the present study may suggest that this population is protected from androgen-driven plasticity in the developmental window that PNA-treatment is applied. Although NPY/AgRP neurons are born on approximately embryonic day 12, significant development in this circuit occurs during the first 3 weeks of the postnatal period (67).

As PCOS-like PNA mice exhibit impaired steroid hormone negative feedback and reduced PR expression in the ARN (14,20), we investigated the steroid hormone sensitivity within NPY\textsuperscript{ARN} neurons specifically. Immunohistochemistry confirmed reduced PR
expressing neurons within the ARN as reported previously (14). Consistent again with this same study, the number of ERα expressing cells in the ARN was unchanged. Despite evidence that NPY expression in the ARN varies over the estrous cycle of both the rat and mouse (68,69), we found almost a complete absence of NPY^{ARN} co-expression with ERα or PR. Kim et al. demonstrate a similarly low co-expression of ERα via IHC in mice while also demonstrating oestradiol reduces NPY mRNA levels (70). Estradiol regulation of NPY via classical receptors is therefore likely indirect through peripheral pathways that then alter NPY expression. The lack of PR co-expression in NPY^{ARN} neurons found here is not surprising given the lack of ERα, and is in line with evidence in the ewe showing that NPY mRNA is not affected by progesterone administration, nor do NPY^{ARN} neurons possess PR (71). These results indicate that the GABA^{ARN} neurons which lose PR expression in PNA mice (14) cannot consist of the NPY expressing subpopulation.

In contrast to ERα and PR, AR was both co-expressed in NPY^{ARN} neurons and up-regulated by PNA treatment. As circulating T levels are elevated in PNA-treated mice (20), and AR expression in the brain appears to be positively auto-regulated by androgens (72,73), it is possible that hyperandrogenism in the PNA mouse is the driver of increased AR expression in NPY^{ARN} neurons. It remains to be determined whether elevated AR impacts neuroendocrine regulation in the PNA-treated mouse. Chronic DHT exposure from 3 weeks of age induces a range of metabolic effects including increased body mass and greater adiposity and neuron-specific knockout of AR protects against DHT-induced metabolic and reproductive impairments (74). Thus, excess androgens may positively regulate NPY^{ARN} neurons to increase orexigenic drive. However, prenatal androgen exposure that leads to postpubertal hyperandrogenism does not increase body weight and results in only very mild metabolic disturbances (53,54), suggesting AR signalling is not elevating NPY activity. Furthermore, as NPY/AgRP appear to inhibit GnRH neurons (38), and activated NPY^{ARN} neurons restrain LH secretion (35), elevated NPY^{ARN} activity would be at odds with the decreased pulse frequency present in PNA-treated mice (20). Therefore, further work is needed to dissect the actions of androgen signalling in NPY^{ARN} neurons and whether these actions play a role in the PCOS-like neuroendocrine impairments associated with prenatal androgen excess.

While NPY^{ARN} innervation to GnRH neurons appears unaltered, it remains possible that alterations to GnRH neuronal afferents are present; for example, NPY^{ARN} neurons project to and regulate kisspeptin neurons in the ARN (39). Closer examination of these afferent
circuits would give a more complete picture of whether PNA alters circuits in the pulse-generating networks of the hypothalamus. While more NPY\textsuperscript{ARN} neurons appear androgen-sensitive, this raises questions regarding the possible functional significance of this increase. As it stands there is very little information in the mouse to suggest how androgens regulate NPY\textsuperscript{ARN} neurons. Future studies will be needed to determine if androgens acting via AR in NPY\textsuperscript{ARN} neurons have any role in the interference of steroid hormone negative feedback within the PNA-treated mouse.

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Figure Legends:

Figure 1. Presence of close appositions between NPY<sup>ARN</sup> neurons and GnRH neurons is region-dependent and not modified by PNA-treatment. (A,B) Representative images composed from z-projections (thickness 12-14 µm) of GnRH neurons (magenta) residing in the MS (i), rPOA (ii) and AHA (iii) of VEH- (A) and PNA-treated (B) mice showing GFP-expressing NPY<sup>ARN</sup> fibers (green) in close contact with GnRH neurons. (C) The mean number of contacts/GnRH neuron made by NPY<sup>ARN</sup> neuron fibers to GnRH neurons located in the MS, rPOA, and AHA, at the level of the soma and the primary dendrite in VEH- (grey bars, n=5) and PNA-treated mice (blue bars, n=8). (D) The mean density of contacts made onto GnRH neurons in the rPOA by ARN NPY fiber projections, at the level of the soma and in 15 µm subsections of the primary dendrite. Results are presented as mean ± SEM. No significant differences were detected by two-way ANOVA (C) or one-way ANOVA (D). Scale bars = 5 µm.

Figure 2. Expression of AR within NPY<sup>ARN</sup> neurons is elevated in PNA-treated mice. (A, top) Representative images composed from z-projections (6.36 µm thickness) collected throughout the ARN of a VEH-treated mouse showing AR-ir (magenta) and τGFP reporter expression of NPY<sup>ARN</sup> neurons (green). (A, bottom) single plane high magnification images taken within the highlighted area indicated in the top row where GFP and AR labelling is observed. (B, top) Representative images composed from 6.36 µm thick z-projections containing AR and GFP-ir in the ARN of a PNA-treated mouse. (B, bottom) single plane high magnification images taken within the highlighted area. (C) Mean number of GFP expressing NPY neurons counted in VEH- (grey bars, n=4) and PNA-treated mice (blue bars, n=4) across the whole ARN. (D) Mean number of AR-ir cells across the entire ARN in VEH- and PNA-treated mice. (D) The percentage of NPY neurons co-expressing AR in VEH- and PNA-treated mice, averaged across the whole ARN. Results are presented as mean ± SEM. * p < 0.05 as determined by Mann-Whitney U test. Scale bars = 100 µm (top), 10 µm (bottom).

Figure 3. Fewer PR-ir cell in the ARN of PNA mice and near complete absence of PR in NPY neurons. (A) Representative single z-plane images from rostral to caudal zones in the ARN showing PR-ir (magenta) and amplified τGFP reporter expression of NPY<sup>ARN</sup> neurons (green) in a VEH-treated mouse (A) and a PNA-treated mouse (B). Inset boxes show enlarged regions of the middle ARN indicating that although PR-ir cells and τGFP expressing NPY<sup>ARN</sup> neurons are in close proximity, there is an almost complete absence of co-
localisation. (C) Mean number of GFP-positive NPY neurons across the whole ARN in VEH- (grey bars, n = 8) and PNA-treated (blue bars, n = 10) mice. (D) Mean number of PR-ir cells across the whole ARN in VEH- and PNA-treated mice. Results are presented as mean ± SEM. **, p < 0.01 as determined by a two-tailed unpaired Student’s t-test. Scale bars = 50 µm, scale bar in inset 10 µm.

Figure 4. ERα co-expression with NPYARN neurons is absent in both VEH- and PNA-treated mice. (A, top) Representative images composed from z-projections (6.36 µm thickness) collected throughout the ARN of a VEH-treated mouse showing ERα-ir (magenta) and τGFP reporter expression of NPYARN neurons (green). (A, bottom) Single plane high magnification images taken within the highlighted area indicated in the top row. Individual single-labelled NPY neuron cell bodies are indicated by the empty arrowheads, while ERα-ir nuclei are shown in close proximity. (B) Mean number of GFP-expressing NPY neurons in VEH- (grey bars, n=5) and PNA-treated (blue bars, n=5) mice across the whole ARN. (C) Mean number of ERα-ir cells counted in VEH- and PNA-treated mice across the whole ARN. Results are presented as mean ± SEM. No significant differences were detected by two-tailed unpaired Student’s t-test. Scale bars = 100 µm (top), 10 µm (bottom).
Table 1. The mean number of steroid hormone receptor-positive and GFP-positive cells in the rostral, middle and caudal ARN of VEH- and PNA-treated mice.

|        | rARN |        | mARN |        | cARN |
|--------|------|--------|------|--------|------|
|        | AR   | GFP    | %    | AR     | GFP  | %    |
| VEH    | 222.6 ± 16.6 | 82.0 ± 5.9 | 22.7 ± 1.6 % | 237.8 ± 23.6 | 88.1 ± 4.0 | 22.3 ± 3.4 % | 138.0 ± 9.27 | 59.1 ± 5.7 | 15.2 ± 2.8 |
| PNA    | 300.5 ± 7.6 ** | 109.9 ± 16.7 | 32.6 ± 6.1 % | 245.4 ± 6.9 | 86.4 ± 6.0 | 32.6 ± 5.1 % | 178.5 ± 21.1 | 50.1 ± 14.2 | 34.4 ± 5.4 |
| p-value| 0.007 | 0.19 | 0.62 | > 0.99 | > 0.99 | 0.57 | 0.25 | > 0.99 | 0.06 |

|        | AR     | GFP    | %    | AR     | GFP   | %    |
|--------|--------|--------|------|--------|-------|------|
|        | ERα    | GFP    | %    | ERα    | GFP   | %    |
| VEH    | 148.7 ± 7.2 | 92.8 ± 6.9 | 0.0 ± 0.0 % | 205.6 ± 8.3 | 103.2 ± 7.5 | 0.0 ± 0.0 % | 159.4 ± 7.3 | 93.7 ± 7.0 | 0.0 ± 0.0 % |
| PNA    | 180.9 ± 9.6 | 101.5 ± 8.7 | 0.12 ± 0.06 % | 210.7 ± 12.1 | 111.4 ± 9.0 | 0.19 ± 0.10 % | 178.1 ± 11.8 | 104.0 ± 8.9 | 0.13 ± 0.07 % |
| p-value| 0.08 | > 0.99 | 0.40 | > 0.99 | > 0.99 | 0.07 | 0.54 | > 0.99 | 0.32 |

|        | AR     | GFP    | %    | AR     | GFP   | %    |
|--------|--------|--------|------|--------|-------|------|
|        | PR     | GFP    | %    | PR     | GFP   | %    |
| VEH    | 123.9 ± 4.9 | 89.6 ± 9.0 | 0.46 ± 0.21 % | 153.7 ± 9.0 | 105.5 ± 11.9 | 0.31 ± 0.16 % | 134.0 ± 12.5 | 91.8 ± 9.5 | 0.36 ± 0.22 % |
| PNA    | 87.0 ± 8.2 * | 95.9 ± 9.1 | 0.35 ± 0.16 % | 121.3 ± 7.6 | 120.8 ± 13.9 | 0.24 ± 0.08 % | 116.9 ± 12.8 | 100.1 ± 11.8 | 0.17 ± 0.07 % |
| p-value| 0.03 | > 0.99 | > 0.99 | 0.07 | > 0.99 | > 0.99 | 0.66 | > 0.99 | > 0.99 |

Results are presented as mean ± SEM. Columns with % report the proportion of GFP-expressing cells co-localized with steroid hormone receptors. * p < 0.05, ** p < 0.01, VEH vs. PNA within ARN region.
Figure 3

A

VEH

mARN

cARN

GFP (AgRP/PR)

B

PNA

C

GFP-ir NPYARN cells

Number of cells

Treatment

VEH

PNA

D

PR-ir cells in ARN

Number of cells

Treatment

VEH

PNA
Figure 4

A

B

C

GFP-ir NP<sup>tyH<sub>4</sub></sup> cells

ERα-ir cells in ARN

Number of cells

Treatment

YEH

PMA

*