Hypothalamic perineuronal nets are regulated by sex and dietary interventions

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20 **ABSTRACT**

21 Perineuronal nets (PNNs) are widely present in the hypothalamus, and are thought to provide physical
 22 protection and ion buffering for neurons, and regulate their synaptic plasticity and intracellular signaling.
 23 Recent evidence indicates that PNNs in the mediobasal hypothalamus plays an important role in the
 24 regulation of glucose homeostasis. However, whether and how hypothalamic PNNs are regulated are
 25 not fully understood. In the present study, we examined whether PNNs in various hypothalamic regions
 26 in mice can be regulated by sex, gonadal hormones, dietary interventions, or their interactions. We
 27 demonstrated that gonadal hormones are required to maintain normal PNNs in the arcuate nucleus of
 28 hypothalamus in both male and female mice. In addition, PNNs in the terete hypothalamic nucleus
 29 display a sexual dimorphism with females higher than males, and high-fat diet feeding increases terete
 30 PNNs only in female mice but not in male mice. On the other hand, PNNs in other hypothalamic regions
 31 are not influenced by sex, gonadal hormones or dietary interventions. In summary, we demonstrated
 32 that hypothalamic PNNs are regulated in a region-specific manner and these results provide a framework
 33 to further investigate the potential functions of PNNs in regulating energy/glucose homeostasis at the
 34 interplay of sex, gonadal hormones and diets.

35

36

37 **Introduction**

38 Obesity is now recognized as a serious global health problem due to its increasing prevalence and
 39 comorbidities, e.g. the metabolic syndrome. The WHO reported that over 650 million adults worldwide
 40 were obese in 2016 and 40 million children under the age of 5 were overweight or obese in 2018. In U.S.,
 41 the prevalence of adult obesity was 42.4% in 2017~2018 according to the Centers for Disease Control
 42 and Prevention. The etiology of human obesity is not fully understood, and effective treatments for
 43 obesity and associated metabolic disorders are limited. Recent studies revealed genetic and epigenetic
 44 basis for variations in human body mass index (BMI) (1-3), and strikingly the majority of BMI-associated
 45 genetic variants affect genes that are enriched in the brain (3,4). In particular, the brain hypothalamus
 46 receives metabolic and/or hormonal signals reflecting the body's nutritional status and coordinates
 47 neuroendocrine and behavioral responses to maintain body weight balance (5-7). Indeed, various
 48 genetic variants associated with human obesity have been demonstrated to cause energy and/or glucose
 49 dysregulations through impairing functions of neurons and/or neurocircuits within the hypothalamus
 50 (8).

51
 52 Perineuronal nets (PNNs) in the brain are condensed glycosaminoglycan-rich extracellular matrix
 53 structures (9). PNNs typically enmesh neurons in defined circuits, and are thought to provide physical
 54 protection and ion buffering for neurons, and regulate their synaptic plasticity and intracellular signaling
 55 (9). PNNs abundantly exist in the forebrain regions, e.g. the cortex and the hippocampus, and the PNNs
 56 levels in these regions can be regulated by animals' experience, and have been implicated in various
 57 neurobiological disorders such as schizophrenia, bipolar disorder, Alzheimer's disease and addictions.
 58 Recent evidence indicates that chemical disruption of PNNs in the mediobasal hypothalamus
 59 significantly blunts the glucose-lowering effects of central action of fibroblast growth factor 1 (FGF1) in

60 obese Zucker diabetic fatty (ZDF) rats (10). Thus, hypothalamic PNNs may also play important roles in
 61 the regulation of energy and/or glucose homeostasis. Indeed, abundant PNNs are present in multiple
 62 hypothalamic regions (11,12), but whether and how these hypothalamic PNNs are regulated are not fully
 63 understood.

64

65 In the present study, we sought to examine whether PNNs in various hypothalamic regions can be
 66 regulated by dietary interventions in mice. Since many hypothalamic regions or neural populations are
 67 sexually dimorphic and/or regulated by gonadal hormones (13,14), we also explored the potential effects
 68 of sex and/or gonadal hormones on PNN levels. Our results revealed that sex, gonadal hormones and/or
 69 dietary interventions can regulate PNNs in a region-specific manner and provide a framework to further
 70 investigate the potential functions of PNNs in regulating energy/glucose homeostasis at the interplay of
 71 sex, gonadal hormones and diets.

72

73 **METHODS**

74 **Study Animals**

75 C57BL6j male and female mice were group-housed respectively until 14 weeks of age in a temperature-
 76 controlled room under a 12h:12h light-dark cycle with ad lib access to regular chow diet (5V5R-Advanced
 77 Protocol PicoLab Select Rodent 50 IF/6F, PicoLab) and water. Forty nine mice are divided into 16 groups
 78 by four factors (1) sex: male or female groups; (2) surgeries: receiving sham or castration
 79 (CAST)/ovariectomy (OVX) surgeries at 14 weeks of age (see below); (3) dietary interventions: chronic
 80 feeding with chow or a high-fat diet (HFD, D12492i Rodent Diet With 60 kcal% Fat, Research Diets, Inc.),
 81 (4) nutritional status: fed or fasted at the time of perfuse (see below). Care of all animals and procedures
 82 were approved by the Baylor College of Medicine Institutional Animal Care and Use Committee.

83

84 **Surgery**

85 At the age of 14 weeks, either bilateral gonadectomy (CAST for males and OVX for females) or sham
 86 surgeries were executed in these mice. The mice were anesthetized with inhaled 2% isoflurane. For OVX
 87 in females, the dorsolateral incisions (2 cm) were made, followed by blunted separation of the
 88 subcutaneous fat, and incision (1.5cm) the abdominal wall to expose the reproductive tract. The ovaries
 89 were isolated and excised. The reproductive tract was returned to the peritoneum and the incisions on
 90 the abdominal wall were sutured using sterile Vicryl thread (size 6); the skin incisions were sutured by
 91 sterile Prolene thread. For sham surgeries in female mice, the procedures were the same, except that
 92 the ovaries were kept intact. For castration in males, a central vertical incision (3 cm) was made, followed
 93 by blunted separation of any subcutaneous fat to locate the vas deferens. The vas deferens were ligated
 94 with Vicryl; the testes were cut and taken away from the fat pad. The skin incisions were sutured by
 95 sterile Prolene thread. For sham surgeries in male mice, the procedures were the same, except that the

96 testes were kept intact.

97

98 **Food intake, body weight, glucose and body composition**

99 One week after the surgeries, mice were randomly divided into two groups to be either regular chow or
100 HFD described above ad libitum for four weeks. Food intake, body weight and fed glucose were
101 measured before the surgeries and once each week after the surgery for five weeks. The glucose was
102 measured using the glucometer via the tail vein; to avoid confounding effects from circadian clock and
103 feeding behavior on the glucose level, we always measured blood glucose in the morning of the day and
104 after a 2-hour short fasting to ensure the empty stomach. Body composition (fat mass and lean mass)
105 was examined with the Bruker minispec mq10 MRS system before the surgeries and 5 weeks afterwards.

106

107 **Perfusion and WFA staining**

108 Five weeks after the surgeries, mice were further divided into fasted and ad libitum groups before the
109 perfusion. For the fasted group, mice were fasted overnight and then perfused in the next morning; for
110 the ad libitum group, both food and water were provided ad libitum at all time before the perfusion. For
111 the perfusion, mice were anaesthetized with inhaled isoflurane and perfused with saline followed by 10%
112 formalin. The brain sections were cut at 25 μ m and collected into five consecutive series. One series is
113 used for the following staining and quantification. Free-floating tissue sections were incubated overnight
114 at room temperature for 22 hours using a 1:500 dilution of biotin-labelled Wisteria floribunda agglutinin
115 (WFA) (L1516, Sigma-Aldrich) in PBS + 0.25% Triton X-100. The sections were then washed and incubated
116 for 2 hours at room temperature in 1:500 Streptavidin protein 488 (21832, Invitrogen). Images were
117 captured using a fluorescence microscope with 1s exposure time and no change of contrast for all
118 pictures to visualize various hypothalamic nuclei, including the arcuate hypothalamic nucleus (ARH), the

terete hypothalamic nucleus (TE) , the paraventricular hypothalamic nucleus (PVH), lateral hypothalamus (LH), the anterior hypothalamus (AH), the ventromedial hypothalamic nucleus (VMH), the dorsomedial hypothalamic nucleus (DMH), and the perifornical area of the anterior hypothalamus (PeFAH). Fluorescence intensity of PNNs for each image was quantified using an established method by the macro plugin “Perineuronal net Intensity Program for the Standardization and Quantification of ECM Analysis” (PIPSQUEAK AI v5.3.9, Rewire Neuro, Inc.) (15).

Statistics

Statistical analysis was done using SPSS (SPSS 22.0.0.0, IBM SPSS Statistics) and GraphPad Prism (GraphPad Prism 8.4.2, Graphpad software, LLC). Multivariate linear regression in SPSS was used to investigate the influence of sex, GDX and chronic dietary intervention on various metabolic parameters (cumulative food intake, body weight change, blood glucose change, fat change and lean change). If sex or GDX was found to have a significant effect on any parameter, the p value for the interaction of sex and GDX was tested. If any two factors were found to have significant effects, the p value for the interaction of these two factors was tested.

For the PNN fluorescence intensity, in addition to the sex, GDX and chronic diet intervention, we introduced one more factor (nutritional status) as mice were perfused either after an overnight fasting or ad libitum. Similarly, we first used the multivariate linear regression analyses in SPSS to examine effects of these 4 factors on PNNs fluorescence intensity in each hypothalamic region. If sex or GDX was found to have a significant effect on any parameter, we then used the two-way ANOVA to examine the interaction of sex and GDX. In case that sex had a significant effect, we used the GraphPad Prism to further examine effects of dietary intervention and nutritional status in each sex separately. $P < 0.05$ was

142 considered statistically significant.

143

144 **Study approval**

145 Care of all animals and procedures were approved by the Baylor College of Medicine Institutional Animal

146 Care and Use Committee.

147

148 **Results**

149 **Distribution of WFA-labelled PNNs in the mouse hypothalamus**

150 We first surveyed the distribution of PNNs in the mouse hypothalamus using WFA immunostaining.
 151 Abundant WFA-labelled PNNs were detected in multiple hypothalamic regions, including the LH, the
 152 ARH, the VMH and the TE (Figure 1). In addition, we also observed modest WFA immunoreactivity in the
 153 AH, the PVH and the DMH (Figure 1). Consistent with a previous report (12), WFA-labelled PNNs were
 154 also observed in a newly identified hypothalamic area, namely the PeFAH (Figure 1A).

155

156 **Effects of sex, gonadal hormones and dietary interventions on metabolic parameters**

157 Given the essential roles of the hypothalamus in regulating energy homeostasis, we sought to examine
 158 potential effects of dietary interventions on the level of PNNs in the hypothalamus. In order to explore
 159 potential influence by sex and/or gonadal hormones, we included both male and female mice with or
 160 without gonadectomy (GDX) in these analyses. Briefly, male and female C57Bl6j mice underwent sham
 161 or GDX (castration for males and ovariectomy for females) at the 14 weeks of age, followed by 5-week
 162 chronic feeding with either a regular chow diet or HFD. As expected, the multivariate linear regression
 163 analysis revealed that compared to chow-fed groups, HFD feeding had profound effects on multiple
 164 metabolic parameters, including significantly increasing calorie intake, blood glucose, body weight gain
 165 and fat mass gain (Table 1). Notably, fat mass gain was significantly smaller in females than in males
 166 (Table 1). We also noted that lean mass gain was significantly larger in females than in males, and that
 167 GDX significantly reduced lean mass gain; in addition, there was a significant interaction between sex
 168 and GDX on this parameter (Table 1). In summary, multiple metabolic parameters in mice were
 169 influenced by sex, gonadal hormones, the dietary interventions, and/or their interactions, as reported
 170 by others (16,17).

171

172 **Effects of gonadal hormones on PNN in the ARH**

173 At the end of 5-week feeding, mice were then perfused either ad libitum or after an overnight fasting.
 174 We then used these fixed brain samples for WFA immunostaining and then quantified the level of WFA-
 175 labelled PNNs in each hypothalamic region in various conditions. In the ARH, the WFA fluorescence
 176 intensity was not significantly affected by chronic HFD feeding (vs. chow), or by the acute fasting (vs. ad
 177 libitum), as revealed by the multivariate linear regression analysis (Table 2). Interestingly, we noted that
 178 GDX significantly reduced WFA fluorescence intensity in the ARH compared to the sham group (Table 2
 179 and Figure 2A-2E). A detailed analysis at different rostral-caudal levels revealed that GDX mainly reduced
 180 WFA in the rostral to medial ARH (-1.94 to -1.58 mm to Bregma) but the caudal ARH was not affected
 181 (Figure 2F). Because female and male mice experienced different GDX surgeries (OVX for females and
 182 CAST for males), we used a two-way ANOVA analysis to further analyze the interaction between sex and
 183 GDX. We found that the inhibitory effect of GDX was independent of sex; in other words, both OVX in
 184 females and CAST in males significantly reduced WFA fluorescence intensity in the ARH (Table 2 and
 185 Figure 2G). We also examined effects of dietary innervations on ARH PNNs in sham or in GDX groups,
 186 respectively, but did not detect any significant effect of by either chronic HFD feeding or by acute fasting
 187 (data not shown). Thus, these results indicate that endogenous gonadal hormones in both males and
 188 females are required to maintain normal PNNs in the ARH.

189

190 **Effects of sex and HFD feeding on PNN in the TE**

191 In the TE, female mice showed significantly higher WFA fluorescence intensity compared to males, as
 192 revealed by the multivariate linear regression analysis (Table 3 and Figure 3A-3E), and this sex difference
 193 existed in a rostral subdivision (-1.70 mm to Bregma) and a medial-caudal subdivision (-2.18 to -2.30 mm

to Bregma) of the TE (Figure 3F). Using a two-way ANOVA analysis, we further found that the significant sex difference only existed in sham mice, but not in GDX mice (Figure 3G). Thus, these results indicate that gonad-intact female mice have higher PNNs in the TE than gonad-intact males.

Given the clear influence by sex, we reanalyzed the effects of HFD feeding in each sex individually and used a two-way ANOVA analysis followed by Sidak tests to further explore potential interactions between chronic HFD feeding and acute fasting. In female mice fed ad libitum, chronic HFD feeding significantly enhanced WFA fluorescence intensity in the TE compared to chow-fed females, but this effect was not observed after an overnight fasting (Figure 4A-4E). This HFD-induced increases existed in a rostral-medial subdivision (-1.70 to -1.94 mm to Bregma) of the TE (Figure 4F). On the other hand, in male mice, neither chronic HFD feeding nor acute fasting induced any changes in WFA fluorescence intensity in the TE (Figure 4G). Thus, these results indicate that chronic HFD feeding can increase PNNs in the TE in female mice but not in male mice.

Other hypothalamic regions

We also analyzed effects of sex, gonadal hormones, and/or dietary interventions on WFA fluorescence intensity in all other hypothalamic regions. The multivariate linear regression analysis revealed that WFA fluorescence intensity in the PVH, the PeFAH, the LH, the AH, the VMH or the DMH, was not significantly altered by these conditions (Table 4).

214 **Discussions**

215 One interesting observation of our study is that endogenous gonadal hormones are required to maintain
 216 normal PNNs in the ARH in both male and female mice. The ARH in the hypothalamus has been long
 217 believed to contain the first order neurons that respond to the peripheral signals, including leptin (18-
 218 20), insulin (21-23), ghrelin (24), asprosin (25) and estrogens (26-28). The ARH neurons include those
 219 expressing pro-opiomelanocortin (POMC) and those co-releasing agouti-related peptide (AgRP),
 220 neuropeptide Y (NPY), and GABA (referred to as AgRP neurons). POMC neurons synthesize and secrete an
 221 anorexigenic peptide, α -melanocyte-stimulating hormone (α -MSH) which prevents overeating and body
 222 weight gain (29,30). On the other hand, AgRP neurons are orexigenic (31), and activation of AgRP
 223 neurons promotes eating even when mice are satiated (32,33). Recent evidence also indicates that non-
 224 AgRP GABAergic neurons in the ARH provide a redundant orexigenic mechanism to drive feeding and
 225 body weight gain (34). Notably, the majority of PNNs in the ARH enmeshes AgRP neurons and GABAergic
 226 neurons, with only a small portion enmeshing POMC neurons (11). Interestingly, reduced PNNs in the
 227 ARH is found in ob/ob mice deficient in the leptin gene and in ZDF rats with a mutated leptin receptor
 228 gene (10,11), highlighting an important role of leptin signaling in maintaining the normal PNNs in the
 229 ARH. Further, the central administration of FGF1 in ZDF rats can enhance PNNs in the ARH, associated
 230 with a prolonged glucose-lowering benefit (10). Importantly, chemical disruption of PNNs significantly
 231 shortens this glucose-lowering effect of FGF1 (10). Thus, PNNs in the ARH play an essential role in
 232 regulating glucose balance.

233
 234 Here we demonstrated that PNNs in the ARH are significantly reduced by removal of the ovaries in
 235 female mice, suggesting that PNNs are also regulated by ovarian hormones. Supporting this notion, co-
 236 administration of both ovarian hormones (estrogen and progesterone), which simulates pregnancy, can

trigger PNN formation in the medial preoptic area, a hypothalamic region normally devoid of PNNs in non-pregnant females (35). While the role of progesterone in energy and glucose homeostasis is not clear (14), estrogen is well established to provide many metabolic benefits, including suppressing food intake, increasing energy expenditure, reducing body weight gain, enhancing insulin sensitivity, and lowering blood glucose (16,36-41). Both estrogen receptor α (ER α) and estrogen receptor β (ER β) are implicated in mediating these metabolic effects (42-44). Notably, abundant ER α is expressed by POMC neurons in the ARH (45-47), to a less extent by AgRP neurons (48,49), while ER β expression in the ARH is minimal (50). We recently showed that intact ER α signals are required for the glucose-sensing functions of ARH neurons (51). Further, estrogen can enhance glutamatergic synapses onto POMC neurons (52), and increase their excitability (53,54). Importantly, we previously showed that selective deletion of ER α from POMC-lineage neurons in female mice results in hyperphagia and obesity (26), and attenuates estrogen-induced anorexia (55). Thus, these findings, together with our current observations that PNNs are reduced by ovariectomy, suggest that PNNs in the ARH may mediate, at least a portion of, estrogenic actions on energy and glucose homeostasis, and future investigations are warranted to test this possibility. Interestingly, we also observed similar reductions of PNNs in the ARH of male mice depleted with endogenous testosterone. Consistently, it has been reported that testosterone administration can stimulate PNN formation in the forebrain of song birds (56). However, it is worth noting that testosterone can be converted into estrogen by aromatase (57). Thus, it remains to be tested whether the testosterone-induced PNN expression is mediated by testosterone per se or by converted estrogen.

Another hypothalamic region where PNNs are regulated is the TE, where female mice showed higher PNNs than males. Importantly, this sex difference was only observed in gonad-intact mice, but was

blunted in mice with gonads surgically removed. These results suggest that PNNs in the TE may be also regulated by gonadal hormones. Supporting this possibility, estrogen has been shown to increase oxytocin binding in the TE in rats (58), indicating potential actions of estrogen in this region. It has to be pointed out that, unlike other hypothalamic regions (e.g. the ARH, VMH, etc.), the TE has received very little attention with no more than 5 publications regarding this nucleus available in the PubMed. Early neuroanatomic studies reported that cell bodies of cholecystokinin neurons and calcitonin gene-related peptide (CGRP) neurons are found in the TE (59,60). In addition, the TE harbors nerve fibers that are immunoreactive for NPY, α -MSH, β -endorphin, corticotropin-releasing hormone, galanin, and substance P (59,60), suggesting that the TE neurons receive a wide range of innervations and inputs from other neural populations. Interestingly, we found that in female mice only, PNNs in the TE was upregulated by chronic HFD feeding, while such effect was not observed after an overnight fasting. It is worth noting that among many hypothalamic regions expressing PNNs, the TE is the only region where PNNs can be regulated by dietary interventions. Considering that the TE receives inputs from NPY and α -MSH, both of which are implicated in the regulation of feeding and body weight, we suggest that TE neurons and the PNNs enmeshing these neurons may play an under-appreciated role in the context of energy homeostasis. This possibility warrants future investigations.

In summary, here we systemically assessed PNNs in multiple hypothalamic regions and explored potential regulations by sex, gonadal hormones, dietary interventions, and their interactions. We acknowledge that the limitation of the current study is its descriptive nature. However, we want to point out that our results filled a gap of knowledge in the relevant field and provided a framework to develop new hypotheses and design future studies to determine the physiological functions of hypothalamic PNNs and to explore the regulatory mechanisms. For example, the strong regulations by gonadal

283 hormones on PNNs in the ARH suggest that gonadal hormones may act through PNNs in the ARH to
 284 regulate energy and/or glucose homeostasis. Further, given that PNNs in the ARH are required to
 285 mediate metabolic benefits of FGF1 (10), the efficacy of FGF1 may need to be examined in animals
 286 depleted of gonadal hormones, which would provide important pre-clinical data for potential
 287 applications of FGF1 or its analogs in aged men with testosterone deficiency or in post-menopausal
 288 women. In addition, the dietary regulations on PNNs in the TE suggest that this understudied
 289 hypothalamic nucleus may play a role in the regulation of feeding and body weight balance. Importantly,
 290 since these dietary regulations only existed in female mice but not in male mice, future studies need to
 291 include at least female animals, if not both. Only including male animals (a common practice in many
 292 laboratories) might miss potential discoveries about PNNs in the TE or TE neurons themselves.
 293

294 **Author Contributions:** NZ was involved in experimental design and most of procedures, data acquisition
 295 and analyses, and writing the manuscript. ZY assisted in data analysis. HL, MY, YH, HL, CL, LT, LW, NY, JH,
 296 NS, YY and CW assisted in in surgical procedures and production of study mice. LLC and TZ are involved
 297 in the study design and manuscript preparation. YX is the guarantors of this work and, as such, had full
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306

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Tables

Table 1. Effects of sex, GDX and diet on various metabolic parameters.

	B	p value	95.0% Confidence Interval	
			Lower Bound	Upper Bound
Cumulative intake (calorie)				
Female vs Male [#]	-13.234	0.298	-38.542	12.075
GDX vs Sham ^{##}	7.461	0.554	-17.798	32.721
High fat diet vs Regular chow ^{###}	56.660	p < 0.001***	31.306	82.013
Body weight change (g)				
Female vs Male [#]	-0.448	0.620	-2.255	1.359
GDX vs Sham ^{##}	0.532	0.556	-1.275	2.339
High fat diet vs Regular chow ^{###}	5.389	p < 0.001***	3.582	7.196
Blood glucose change (mg/dL)				
Female vs Male [#]	13.141	0.128	-3.915	30.198
GDX vs Sham ^{##}	1.143	0.893	-15.913	18.200
High fat diet vs Regular chow ^{###}	37.838	p < 0.001***	20.782	54.894
Fat change (g)				
Female vs Male [#]	-2.052	0.012*	-3.633	-0.471
GDX vs Sham ^{##}	1.495	0.063	-0.085	3.076
P for the interaction of Sex and GDX		0.451		
High fat diet vs Regular Chow ^{###}	5.539	p < 0.001***	3.958	7.120
P for the interaction of Sex and Diet		0.226		
Lean change (g)				
Female vs Male [#]	1.405	p < 0.001***	0.828	1.983
GDX vs Sham ^{##}	-0.805	0.007**	-1.383	-0.228
P for the interaction of Sex and GDX		p < 0.001***		
High fat diet vs Regular chow ^{###}	-0.107	0.710	-0.684	0.470

Here we use male mice as reference. ## Here we use sham mice as reference. ### Here we use regular chow as reference.

607 **Table 2. Effects of sex, GDX, diet and fasting on PNNs in the ARH.**

	B	p value	95.0% Confidence Interval	
			Lower Bound	Upper Bound
Female vs Male#	3216.040	0.710	-14091.082	20523.162
GDX vs Sham##	-28525.814	0.002**	-45832.935	-11218.692
High fat diet vs Regular chow###	5478.986	0.527	-11858.261	22816.234
Fast vs Ab libitum####	1070.635	0.902	-16266.612	18407.883

608 # Here we use male mice as reference. ## Here we use sham mice as reference. ### Here we use
609 regular chow as reference. #### Here we use fed mice as reference.

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613 **Table 3. Effects of sex, GDX, diet and nutritional status on PNNs in the TE.**

	B	p value	95.0% Confidence Interval	
			Lower Bound	Upper Bound
Female vs Male [#]	37265.520	0.004**	12278.584	62252.456
GDX vs Sham ^{##}	8878.365	0.478	-16108.572	33865.301
High fat diet vs Regular chow ^{###}	3443.149	0.783	-21587.281	28473.579
Fast vs Ab libitum ^{####}	-2328.269	0.852	-27358.698	22702.161

614 # Here we use male mice as reference. ## Here we use sham mice as reference. ### Here we use regular
615 chow as reference. #### Here we use fed mice as reference.

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618 **Table 4. Effects of sex, GDX, diet and nutritional status on PNNs in the PVH, PeFAH, LH, AH, VMH and**
619 **DMH.**

		B	p value	95.0% Confidence Interval	
				Lower Bound	Upper Bound
PVH					
	Female vs Male [#]	1093.487	0.785	-6950.901	9137.874
	GDX vs Sham ^{##}	1801.150	0.649	-6123.285	9725.586
	High fat diet vs Regular chow ^{###}	7111.050	0.082	-948.109	15170.209
	Fast vs Ab libitum ^{####}	1437.498	0.721	-6621.661	9496.657
PeFAH					
	Female vs Male [#]	10127.575	0.447	-16501.677	36756.827
	GDX vs Sham ^{##}	12003.295	0.361	-14228.883	38235.473
	High fat diet vs Regular chow ^{###}	12968.496	0.332	-13709.655	39646.646
	Fast vs Ab libitum ^{####}	3879.818	0.771	-22798.333	30557.969
LH					
	Female vs Male [#]	48723.960	0.118	-12839.004	110286.924
	GDX vs Sham ^{##}	25737.089	0.397	-34907.897	86382.074
	High fat diet vs Regular chow ^{###}	-22043.519	0.475	-83719.530	39632.491
	Fast vs Ab libitum ^{####}	-17950.520	0.560	-79626.531	43725.490
AH					
	Female vs Male [#]	9830.509	0.347	-11017.746	30678.764
	GDX vs Sham ^{##}	6991.586	0.496	-13545.796	27528.969
	High fat diet vs Regular chow ^{###}	6079.521	0.560	-14807.017	26966.059
	Fast vs Ab libitum ^{####}	599.040	0.954	-20287.498	21485.578
VMH					
	Female vs Male [#]	8833.935	0.639	-28921.823	46589.693
	GDX vs Sham ^{##}	6705.915	0.718	-30486.859	43898.688

	High fat diet vs Regular chow ^{###}	-14866.204	0.432	-52691.292	22958.884
	Fast vs Ab libitum ^{####}	-19095.426	0.314	-56920.514	18729.662
DMH					
	Female vs Male [#]	-2075.195	0.135	-4826.375	675.985
	GDX vs Sham ^{##}	267.179	0.843	-2442.978	2977.336
	High fat diet vs Regular chow ^{###}	1587.169	0.252	-1169.064	4343.401
	Fast vs Ab libitum ^{####}	-487.298	0.723	-3243.531	2268.934

Here we use male mice as reference. ## Here we use sham mice as reference. ### Here we use regular chow as reference. #### Here we use fed mice as reference.

Figure Legends

Figure 1. Distribution of WFA-labelled PNNs in mouse hypothalamus. Representative fluorescent microscopic images showing WFA-labelled PNNs (green) and DAPI counterstaining (purple) in a series of coronal brain sections at the level of -0.82 mm (A), -1.34 mm (B), -1.58 mm (C) and -1.82 mm (D) relative to the Bregma. Scale bars = 200 μ m. AH, the anterior hypothalamus; ARH, the arcuate hypothalamic nucleus; DMH, the dorsomedial hypothalamic nucleus; LH, the lateral hypothalamus; PeFAH, the perifornical area of the anterior hypothalamus (PeFAH); PVH, the paraventricular hypothalamic nucleus; TE, the terete hypothalamic nucleus; VMH, the ventromedial hypothalamic nucleus.

Figure 2. GDX reduces PNNs in the ARH. (A-D) Representative fluorescence microscopic images showing WFA-labelled PNNs (green) in the ARH of female sham (A), female OVX (B), male sham (C) and male CAST mice (D). Scale bars = 200 μ m. ARH, the arcuate hypothalamic nucleus. (E) Quantification of the total WFA fluorescence intensity in the ARH from sham vs. GDX mice. N = 24 or 25 mice per group. Data are presented with mean $\pm$ SEM with individual data points. **, P<0.01 in unpaired two-tailed Student's t-tests. (F) Quantification of the WFA fluorescence intensity at various rostral-caudal levels of the ARH from sham vs. GDX mice. N = 24 or 25 mice per group. Data are presented with mean $\pm$ SEM. *, P<0.05 and **, P<0.01 in unpaired two-tailed Student's t-tests at each level. (G) Quantification of the total WFA fluorescence intensity in the ARH from female or male mice with sham or GDX surgeries. N = 12 or 13 mice per group. Data are presented with mean $\pm$ SEM with individual data points. *, P<0.05 in two-way ANOVA analysis followed by Holm-Sidak post hoc tests.

645 **Figure 3. PNNs in the TE display a sexual dimorphism.** (A-D) Representative fluorescence microscopic
 646 images showing WFA-labelled PNNs (green) in the TE of female sham (A), male sham (B), female OVX
 647 (C), and male CAST mice (D). Scale bars = 200 μ m. TE, the terete hypothalamic nucleus. (E) Quantification
 648 of the total WFA fluorescence intensity in the TE from female vs. male mice. N = 24 or 25 mice per group.
 649 Data are presented with mean $\pm$ SEM with individual data points. **, P<0.01 in unpaired two-tailed
 650 Student's t-tests. (F) Quantification of the WFA fluorescence intensity at various rostral-caudal levels of
 651 the TE from female vs. male mice. N = 24 or 25 mice per group. Data are presented with mean $\pm$ SEM. *,
 652 P<0.05 and **, P<0.01 in unpaired two-tailed Student's t-tests at each level. (G) Quantification of the
 653 total WFA fluorescence intensity in the TE from female or male mice with sham or GDX surgeries. N = 12
 654 or 13 mice per group. Data are presented with mean $\pm$ SEM with individual data points. **, P<0.01 in
 655 two way ANOVA analysis followed by Sidak post hoc tests.

656

657 **Figure 4. HFD feeding increases TE PNNs in female mice.** (A-D) Representative fluorescence microscopic
 658 images showing WFA-labelled PNNs (green) in the TE of female mice that were fed chow ad libitum (A),
 659 HFD ad libitum sham (B), chow after an overnight fasting (C), and HFD after an overnight fasting (D).
 660 Scale bars = 200 μ m. TE, the terete hypothalamic nucleus. (E) Quantification of the total WFA
 661 fluorescence intensity in the TE from female mice with various dietary interventions. N = 6 mice per
 662 group. Data are presented with mean $\pm$ SEM with individual data points. *, P<0.05 in two-way ANOVA
 663 analysis followed by Sidak post hoc tests. (F) Quantification of the WFA fluorescence intensity at various
 664 rostral-caudal levels of the TE from female mice fed with chow or HFD ad libitum. N = 6 mice per group.
 665 Data are presented with mean $\pm$ SEM. *, P<0.05 and **, P<0.01 in unpaired two-tailed Student's t-tests
 666 at each level. (G) Quantification of the total WFA fluorescence intensity in the TE from male mice with

667 various dietary interventions. N = 6 or 7 mice per group. Data are presented with mean $\pm$ SEM with
668 individual data points. No significance was detected by in two way ANOVA analysis.

Figure 1

Figure 2

Figure 3

Figure 4