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Pituitary corticotroph somatostatin receptor subtype 5 (SSTR5) signals to inhibit adrenocorticotrophin (ACTH) secretion. As ACTH deficiency results in attenuated adrenal cortisol production and an impaired stress response, we sought to clarify the role of SSTR5 in modifying the hypothalamic/pituitary/adrenal (HPA) axis. We generated Tg HP5 mice overexpressing SSTR5 in pituitary corticotrophs that produce the ACTH precursor proopiomelanocortin (POMC). Basal ACTH and corticosterone were similar in HP5 and WT mice, while HP5 mice showed attenuated ACTH and corticosterone responses to corticotrophin releasing hormone (CRH). HP5 mice exhibited attenuated corticosterone responses upon a restraint stress test and inflammatory stress following LPS injection, as well as increased anxiety-like and depressive-like behavior on open field and forced swim tests. Pituitary corticotroph CRH receptor subtype 1 (CRHR1) mRNA expression and ACTH responses to CRH were also attenuated in HP5 mice. In AtT20 cells stably overexpressing SSTR5, CRHR1 expression and cAMP response to CRH were reduced, whereas both were increased after SSTR5 KO. In elucidating mechanisms for these observations, we show that SSTR5-induced miR-449c suppresses both CRHR1 expression and function. We conclude that corticotroph SSTR5 attenuates HPA axis responses via CRHR1 downregulation, suggesting a role for SSTR5 in the pathogenesis of secondary adrenal insufficiency.

Introduction

The hypothalamic/pituitary/adrenal (HPA) axis plays a central role in regulating homeostasis, including metabolism, energy storage and expenditure, immunity, and brain functions (1). The HPA axis also enables appropriate stress response. Corticotrophin releasing hormone (CRH) secreted by hypothalamic neurons induces pituitary corticotroph production of the proopiomelanocortin (POMC) precursor polypeptide and enables POMC cleavage by prohormone convertase 1/3 (PC1/3) to produce adrenocorticotrophin (ACTH). Circulating ACTH, in turn, stimulates adrenal zona fasciculata cells to produce cortisol in humans or corticosterone in rodents (1–4). Low circulating levels of cortisol and/or corticosterone, which characterize hypocortisolism, are observed in ACTH-dependent adrenal insufficiency (AI).

Primary AI results from adrenal-related damage, while secondary AI, resulting from pituitary ACTH deficiency, is typically caused by a hypothalamo-hypophyseal lesion or from a central congenital abnormality. Idiopathic secondary AI and tertiary AI resulting from hypothalamic CRH deficiency are rare (5–7). Clinical features of AI include extreme physical and mental fatigue, depression and anxiety, irritability, lack of appetite and weight loss, body pain, nausea and vomiting, diarrhea, fever, hypotension, and hyponatremia (8, 9). If left untreated, especially during stress, AI progresses to adrenal crisis with cardiovascular collapse, often resulting in death (10).

Few animal models are available for the study of secondary AI (11). KO of pituitary developmental transcription factors LHX3 and LHX4, as well as corticotroph transcription factor TPIT, result in corticotroph cell lineage hypoplasia with disrupted ACTH production; these mice have severe baseline hypocortisolism and develop secondary AI (11–13). In KO mice lacking CRH (14, 15) or CRH receptor subtype 1 (CRHR1) (16), ACTH levels at baseline are normal but are attenuated in response to stress. Baseline corticosterone levels are also reduced (14–16).
Somatostatin plays a key role in physiological regulation of the HPA axis (17, 18). Hypothalamic somatostatin inhibits pituitary ACTH secretion (19, 20), mediated mainly by corticotroph somatostatin receptors subtype 2 (SSTR2) and subtype 5 (SSTR5) (21–23). SSTRs, G-protein coupled receptors (GPCRs), inhibit adenylate cyclase, reduce intracellular cAMP levels, close calcium channels to reduce intracellular calcium levels, and acutely inhibit exocytosis (24–26). Pasireotide, a somatostatin receptor ligand with a preferential high affinity for SSTR5, suppresses ACTH and cortisol levels in some patients with Cushing disease caused by ACTH-secreting pituitary adenomas (27, 28).

Consistent with the inhibitory effect of SSTR5 on ACTH secretion, we showed that constitutive human SSTR5 (hSSTR5) activity also inhibits CRH-induced ACTH (29). This action was associated with reduced CRHR1 mRNA expression in mouse AtT20 corticotroph cells overexpressing hSSTR5 (29–31). By contrast, mice devoid of SSTR5 showed increased basal ACTH and corticosterone levels (32). We therefore hypothesized that increased mouse pituitary corticotroph SSTR5 would decrease pituitary ACTH production due to CRHR1 downregulation, resulting in secondary AI and functional decompensation.

We examined the role of corticotroph SSTR5 in adaptive HPA axis responses to stress and elucidated mechanisms underlying SSTR5 modification of adrenal function. Our results show that corticotroph SSTR5 posttranscriptionally suppresses CRHR1 expression and function via induced microRNA, resulting in attenuated corticotroph responses to hypothalamic CRH.

Results

Generation of human pituitary SSTR5 Tg mice (HP5). Human and mouse SSTR5 share 81% homology (33, 34), and both signal to reduce intracellular calcium and cAMP levels (35). Accordingly, we generated corticotroph-targeted Tg mice expressing pituitary hSSTR5, termed HP5, using a corticotroph-specific artificial promoter (36) driving N-terminally HA-linked hSSTR5 (Figure 1A). The promoter was originally derived from Tg mice employing the rat Pmoc (rPmoc) promoter (37, 38). To enhance corticotroph hSSTR5 specificity and expression levels (39), multiple copies of NeuroD1 and Tpit/Pitx binding sequences were inserted upstream of the rPmoc promoter. Both Western blotting and quantitative PCR (qPCR) showed HA-hSSTR5 expression in the pituitary but not in hypothalamic or adrenal tissue (Figure 1B, Supplemental Figure 1A; supplemental material available online with this article; https://doi.org/10.1172/jci.insight.122932DS1). Female and male HP5 mice demonstrated similar patterns of hSSTR5 expression (data not shown). As female nulliparous mice have higher corticosterone levels than males (40, 41), subsequent experiments were performed with female HP5 mice aged 6–12 months.

To examine pituitary gland distribution of hSSTR5, we dissected the anterior lobe (AL) from the intermediate lobe (IL) and posterior lobe (PL), confirmed by Western blot using pituitary-specific markers (42). As expected, PC1/3 was present in both AL and IL+PL, while PC2 and AVP were expressed in IL+PL; POMC was more abundant in IL+PL (Supplemental Figure 1B). Avp mRNA was restricted to IL+PL, as assessed by qPCR (Supplemental Figure 1C). hSSTR5 was more abundant in IL+PL compared with AL (Supplemental Figure 1, D and E), but it was clearly expressed in AL when separately compared against WT (Supplemental Figure 1F). We observed POMC immunofluorescence in AL and IL in both WT and HP5 pituitary (Figure 1, C and D). hSSTR5 was more abundant in IL compared with AL in HP5 and was not expressed in WT (Figure 1, C and D). Colocalization of hSSTR5 and POMC in HP5 AL was confirmed by confocal immunofluorescence microscopy (Figure 1E), demonstrating corticotroph hSSTR5 expression.

HP5 mice maintain baseline pituitary-adrenal function. In HP5 and WT mice, body weight and food consumption were similar up to 23 months of age (Supplemental Figure 2, A and B). At baseline, morning circulating ACTH, αMSH, and corticosterone levels were similar in HP5 and WT mice (Figure 2, A–C), as were levels of prolactin, insulin-like growth factor–I (IGF-I), fasting serum glucose, and insulin (Supplemental Figure 2, C–F). HP5 adrenal gland size (Figure 2D) and weight (P = 0.004; Figure 2E) were lower than those encountered in WT mice. Accordingly, on H&E staining, adrenal cortex width was narrower in HP5 than in WT mice (0.45 ± 0.02 vs. 0.29 ± 0.03 mm, respectively; P = 0.0009) (Figure 2, F–H).

Attenuated HPA axis response to stress in HP5 mice. To assess HPA axis function in HP5 mice, we measured ACTH and corticosterone before and 30 minutes after a single injection of 40 μg/kg CRH. Although baseline ACTH and corticosterone levels were similar in WT and HP5 mice (Figure 2, A and C), evoked ACTH and corticosterone responses to CRH were attenuated in HP5 mice (ACTH,
$P = 0.06$; Figure 3A; corticosterone, $P < 0.0001$; Figure 3B). Next, we treated mice with the somatostatin receptor ligand pasireotide long-acting release (LAR) in expectation of enhancing hSSTR5 signaling, given its high affinity for SSTR5. After 35 days of pasireotide LAR treatment (Supplemental Figure 3A), IGF-I levels were reduced in both WT ($P = 0.01$) and HP5 mice ($P < 0.0001$) (Supplemental Figure 3D), while unstimulated ACTH and corticosterone levels were not altered (Supplemental Figure 3, B and C). However, pasireotide-treated HP5 mice exhibited a blunted response to CRH, exhibiting
lower ACTH (P = 0.004) and lower corticosterone (P = 0.001) after CRH stimulation compared with pasireotide-treated WT mice (Supplemental Figure 3, E and F).

To evaluate changes in corticotroph SSTR5 signaling under stress conditions, mice were restrained for 30 minutes (physiological stress) or treated with LPS (inflammatory stress). The corticosterone response was attenuated in HP5 vs. WT mice after 30 minutes of restraint (P = 0.008; Figure 3C), as well as 2 hours after 10 mg/kg LPS injection (P = 0.03; Figure 3D). Seventy-two hours after 5 mg/kg LPS injection, HP5 mice lost significant body weight, while WT body weight was not altered (P = 0.02; Figure 3E). Importantly, although WT mice were highly mobile and active with unchanged fur, HP5 mice had lost weight and fur, were emaciated, and were less mobile (Supplemental Video 1). We scored mouse body condition and well-being 72 hours after 10 mg/kg LPS administration (n = 10 per group) and found that more HP5 than WT mice exhibited a body condition score of 1 (BCS1), indicating the poorest health condition (30% vs. 10%). 4 HP5 mice and 1 WT mouse died after LPS injection (Figure 3F). These results demonstrate that HP5 mice tolerated extreme inflammatory response less favorably than did WT mice.

Anxiety- and depressive-like phenotype in HP5 mice. We next assessed HP5 and WT mice for behavioral changes including anxiety and depression, known to be associated with HPA axis dysfunction (43, 44). During a 60-minute open field test, HP5 mice exhibited less locomotor activity (P = 0.05; Figure 4A),
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longer resting time ($P = 0.02$; Figure 4B), and less moving distance ($P = 0.04$; Figure 4C) compared with WT mice, but no difference in traveling speed was seen (Figure 4D). Less rearing behavior was observed in HP5 mice during the first 10 minutes when exploring the new chamber ($P = 0.02$ for 0–5 min; Figure 4E) but not later. HP5 mice also spent less time ($P = 0.04$; Figure 4F) and were less active ($P = 0.05$; Figure 4G) in the center of the cage. Together, these behaviors indicate that HP5 are more anxious than WT mice.

Figure 3. HP5 exhibit attenuated ACTH-corticosterone responses to stress. (A and B) Morning ACTH and corticosterone levels measured before (baseline) and 30 minutes after 40 mg/kg i.p. CRH administration in WT and HP5 mice. (A) ACTH measured in WT ($n = 10$) and HP5 ($n = 9$) mice. (B) Corticosterone measured in WT ($n = 9$) and HP5 ($n = 9$) mice. (C) Morning corticosterone levels before (baseline) and 30 minutes after restraint stress in WT ($n = 10$) and HP5 ($n = 8$) mice. (D) Morning corticosterone levels before (baseline) and 120 minutes after 10 mg/kg LPS injection in WT ($n = 10$) and HP5 ($n = 10$) mice. (E) Body weight before (baseline) and 72 hours after 5 mg/kg LPS injection in WT ($n = 6$) and HP5 ($n = 6$) mice. (F) Death and body condition scores (BCS) 72 hours after 10 mg/kg LPS injection in WT ($n = 10$) and HP5 ($n = 10$) mice. BCS1, emaciated; BCS2, under-conditioned; BCS3, well-conditioned. All experiments were performed with different mouse groups. Results are presented as mean ± SEM. * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$; 2-tailed, unpaired t test.

Pituitary CRHR1 expression and action are attenuated in HP5 mice. We next examined mechanisms underlying SSTR5-mediated attenuation of the HPA axis. Hypothalamic Crh and Avp mRNA expression levels were not altered in HP5 or WT (Figure 5, A and B). However, HP5 pituitary glands expressed significantly less Crhr1 mRNA ($P = 0.001$; Figure 5C) and more Agrp mRNA ($P = 0.05$, Figure 5D), while Tpit (Figure 5E), Pome (Figure 5F), and Pitx1 (Figure 5G) mRNA expression levels were similar to those in WT. Growth hormone (Gh1), serving as a control, was unchanged, indicating a cell-specific corticotroph effect (Figure 5H). Western blot analysis confirmed that whole pituitary CRHR1 abundance

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in HP5 was reduced compared with WT \((P = 0.02; \) Figure 5, I and J), and no difference in TPIT, POMC, and GH expression abundance were noted (Figure 5, I and K–M). Pituitary AL was dissected from IL and PL and analyzed separately by Western blot. AL POMC \((P = 0.004; \) Supplemental Figure 4, A and B) and CRHR1 \((P = 0.01; \) Supplemental Figure 4, A and C) expression were decreased as compared with WT, but AL TPIT and GH expression were similar (Supplemental Figure 4, A, D, and E). In line with these observations, in whole pituitary–derived cell cultures, CRH dose-dependent induction of ACTH response was attenuated in HP5 cells (Figure 5N). In contrast, AVP dose-dependent induction of ACTH response was enhanced (Figure 5O) as compared with WT, suggesting that HP5 mice compensate for reduced CRHR1 levels by increasing the AVP response.

**SSTR5 suppresses corticotroph CRHR1 expression and signaling.** To determine whether SSTR5 alters CRHR1 expression, we treated primary pituitary cells derived from SSTR2-KO mice with pasireotide, which has a high affinity for SSTR5, and employed octreotide, which has a high affinity for SSTR2, as a negative control. Pasireotide but not octreotide suppressed \(Chrh1\) mRNA levels \((P = 0.03; \) Figure 6A), while levels of \(Pitx1, Tpit, NeuroD1,\) and \(Pomc\) mRNA levels were similar (Figure 6A). Furthermore, \(Chrh1\) mRNA \((P = 0.0007)\) and CRHR1 protein \((P = 0.003)\) levels were markedly suppressed in atT20 mouse corticotroph cells stably expressing hSSTR5 compared with control (Figure 6, B–D). To confirm the requirement of SSTR5

![Figure 4. Anxiety- and depressive-like phenotype in HPS mice.](image-url)

(A–G) Exploratory/anxiety behavior in WT \((n = 20)\) and HP5 \((n = 20)\) mice assessed by open-field test. Locomotor activity was assessed by (A) number of horizontal beam breaks using the lower ring of photobeams in the entire chamber; (B) resting time (seconds); (C) distance moved (meters); (D) speed (centimeters/second); (E) rearing, defined as the number of vertical beam breaks using the upper ring of photobeams during each 5-minute time block from 0-10 minutes; (F) center time, defined as the percentage of time spent in the center area; and (G) center activity, defined as the percentage of horizontal beam breaks occurring in the center area. (H–K) Depression-like behavior in WT \((n = 20)\) and HP5 \((n = 20)\) mice assessed by 6-minute forced swim test assessed by (H) duration of swimming (seconds); (I) duration of immobility (seconds); (J) ratio of immobility to total time; and (K) latency time to immobility (seconds). Results are presented as mean ± SEM. *\(P \leq 0.05; **P \leq 0.01; ***P \leq 0.001; 2\)-tailed, unpaired t test.
for CRHR1 suppression, we depleted CRHR1 transfectants by generating 3 independent clones (KO#1, KO#2, and KO#3) of SST5-KO AtT20 cells using CRISPR/Cas9 (Supplemental Figure 5A). Three out-of-frame mutations were confirmed by Sanger sequencing (Supplemental Figure 5, B and C) and predicted defective SST5 transmembrane domains by transmembrane helix prediction program (TMHMM; http://www.cbs.dtu.dk/services/TMHMM/) (Supplemental Figure 5D). Unlike WT cells, which were responsive to the selective SST5 agonist BIM-23206, SST5-KO clones lost responsiveness to BIM-23206, which did not inhibit intracellular cAMP accumulation (Supplemental Figure 5E). These results confirm functional elimination of SST5-related cAMP signaling in these cells. Chr1 mRNA (P = 0.0002, Figure 6E) and

Figure 5. Pituitary CRHR1 expression and action are attenuated in HP5 mice. Hypothalamic expression of (A) Chth and (B) Avp mRNA levels derived from WT (n = 6) and HP5 (n = 6) mice. Relative mRNA levels of (C) Chth1, (D) Avp1b, (E) Tpit, (F) Pomc, (G) Ptx1, and (H) Chth in pituitary tissue from WT (n = 5) and HP5 (n = 5) mice as determined by qPCR and normalized to Gapdh. (I) Western blot analysis of 4 WT and 4 HP5 whole pituitaries for CRHR1, Tpit, Pomc, and GH protein and compared with β-tubulin. (J–M) Quantitative analysis of Western blot results in I performed by scanning densitometry. Protein levels were normalized to β-tubulin. (N) CRH-stimulated ACTH levels measured by ELISA in the culture medium of whole pituitary glands (8.0 × 10^4 cells/well, n = 4 at each CRH concentration) collected from WT (n = 5) and HP5 (n = 5) mice. ACTH level (pg/ml) was normalized to cell number (per 1,000 cells). (O) AVP-stimulated ACTH levels measured by ELISA in the culture medium of whole pituitary glands (2.0 × 10^5 cells/well, n = 4 at each CRH concentration) collected from WT (n = 10) and HP5 (n = 10) mice. ACTH level (pg/ml) was normalized to WST-1 absorbance. Results are presented as mean ± SEM. Experiments were repeated in triplicate (N) or duplicate (O), and cells were plated in quadruplicate wells. Representative results are presented. *P ≤ 0.05; **P ≤ 0.01; ***P ≤ 0.001; 2-tailed unpaired t test, (N and O) with Bonferroni correction.
CRHR1 protein \((P = 0.001; \text{Figure } 6, \text{F and G})\) levels were higher in all SSTR5-KO clones compared with WT controls, suggesting that constitutive SSTR5 activity suppresses CRHR1 expression. By contrast, *P* ≤ 0.05; **P* ≤ 0.01; ***P* ≤ 0.001; ****P* ≤ 0.0001; 2-tailed unpaired t test, (*H* and *I*) with Bonferroni correction.

To test whether SSTR5 negatively regulates CRHR1 downstream signaling, we measured CRH-dependent intracellular cAMP production. Cells overexpressing hSSTR5 exhibited blunted CRH-induced cAMP accumulation compared with controls (Figure 6H), while SSTR5-KO cells showed enhanced cAMP responses to CRH compared with WT controls (Figure 6I). We therefore conclude that SSTR5 suppresses CRH-dependent cAMP production via downregulation of CRHR1. In turn, ACTH production is suppressed, resulting in lower corticosterone production.

**SSTR5 downregulates CRHR1 signaling via miRNA-449c.** To address mechanisms for SSTR5 suppression of *Crhr1* mRNA expression, we first investigated transcriptional regulation of *Crhr1* by SSTR5. Based on data available for human *CRHR1* (45), we studied a region –2.4 kb upstream of the end of the 3′UTR. Fourteen regions were highly conserved between human and mouse (Supplemental Figure 6A), and non-coding sequences of the *Crhr1* locus were selected. ChIP analysis measuring histone acetylation level in these regions showed that a region located in the proximal 5′UTR was highly acetylated compared with the other regions (Supplemental Figure 6B). *Crhr1* promoter activity was therefore analyzed in and adjacent to
this region. We generated constructs to measure mouse Crhr1 promoter activity by luciferase assay (Supplemental Figure 6C) and transfected them into SSTR5-overexpressing AtT20 cells (Tg, Supplemental Figure 6D) or from SSTR5-KO AtT20 cells (KO#1–3, Supplemental Figure 6E). Compared with their respective controls, SSTR5 expression status did not alter Crhr1 promoter activity (Supplemental Figure 6, D and E).

Based on our results, we hypothesized that SSTR5 did not affect Crhr1 transcription but rather reduced mRNA stability, and we therefore explored whether microRNA changes could account for these effects. To test this hypothesis, we identified 25 candidate miRNAs that likely bind mouse Crhr1 3’UTR based on the microRNA database (www.microRNA.org) and chose those with Ct value <35 by qPCR in AtT20 cells and mouse pituitary tissue (data not shown). Twelve miRNAs were suppressed in SSTR5-KO cells ≤0.5-fold in at least 1 of 2 clones compared with WT (Figure 7A and Supplemental Figure 7A). Eight miRNAs were induced ≥1.8-fold in hSSTR5-overexpressing transfectants compared with control (Figure 7A and Supplemental Figure 7B), and expression levels of 8 miRNAs were induced ≥1.8 times in HP5 mouse pituitary compared with control (Figure 7A and Supplemental Figure 7C).

We focused subsequent experiments on 7 miRNAs that showed similar changes in HP5 vs. WT pituitaries, as well as in Tg vs. control AtT20 cells, and that showed contrasting changes in KO and WT AtT20 cells (Figure 7A). We generated oligonucleotide inhibitors to the selected miRNAs, transfected AtT20 WT cells with each of these respective inhibitors, and tested Crhr1 mRNA expression as compared with scramble transfected cells. Inhibition of miR-449c (P = 0.008) and miR-322 (P = 0.03), but not the other tested miRNAs, selectively enhanced Crhr1 expression as assessed by qPCR (Figure 7B). Treatment with a pharmacologic miR-449c inhibitor significantly enhanced CRHR1 protein levels in WT AtT20 cells (P = 0.02; Figure 7C). Similar to reports in other tissues (46), we found that miR-449c also suppresses Myc mRNA in HP5 pituitary (P = 0.006; Supplemental Figure 8A) and that AtT20 cells treated with miR-449c inhibitor showed increased c-Myc protein expression (P = 0.04; Supplemental Figure 8, B and C). Inhibition of miR-449c in primary HP5 pituitary cell cultures rescued CRH-dependent ACTH secretion to the culture medium (Figure 7D). In addition, when we overexpressed miR-449c in AtT20 cells (Figure 7E), only Crhr1 but not other pituitary-specific genes were suppressed (Figure 7F). To confirm that miR-449c participates in SSTR5-mediated CRHR1 inhibition, we transfected miR-449c into SSTR5-KO AtT20 cells and observed marked suppression of CRH-induced cAMP accumulation in the absence of SSTR5 (Figure 7G). Taken together, these results demonstrate that miR-449c likely mediates negative pituitary regulation of CRHR1 by SSTR5 signaling.

Discussion

The HP5 mouse overexpressing pituitary corticotroph SSTR5 recapitulates some clinical and biochemical phenotypes of secondary AI when exposed to behavioral or inflammatory stress. We showed that SSTR5 suppresses CRH action in response to stress by reducing corticotroph cell CRHR1 expression. To our knowledge, this mechanism, whereby a receptor not directly associated with the HPA axis regulates HPA axis function, has not been previously reported.

HP5 and CRHR1-KO mice exhibit some similar features. Adrenal cortex is atrophied, and ACTH and corticosterone levels are attenuated, upon stress in both models (16, 47–49). However, low baseline corticosterone levels are observed in CRHR1-KO but not in HP5 mice as compared with controls (16, 48). Baseline corticosterone and ACTH level may be preserved in HP5 mice due to increased AVP action, as HP5 pituitaries demonstrated higher levels of Avpr1b receptor mRNA and increased response to AVP. Basal AVP levels are also significantly increased in CRHR1-KO mice (48). Thus, our findings suggest that AVP upregulation may compensate for baseline ACTH deficiency. Consistent with normal corticosterone levels in HP5, treatment with pasireotide, a somatostatin receptor ligand with a high affinity for SSTR5, had no further suppressive effect on unstimulated corticosterone levels both in WT and HP5. These results also suggest that SSTR5 may not play a role in basal corticosterone regulation.

Utility of the HP5 mouse is clearly demonstrated when considering the clinical picture of patients with secondary AI. As is seen in CRH-KO and CRHR1-KO mice (14, 16), HP5 mice show deconditioning and significantly attenuated ACTH and corticosterone responses to restraint and inflammatory stress. Importantly, however, HP5 mice show normal growth and development in unstressed conditions and only exhibit significant deterioration in health status when exposed to inflammatory stress. This is akin to human subjects with secondary AI, who show baseline cortisol levels within the low–normal range and, only upon stimulation, demonstrate a lack of appropriate cortisol response. Indeed, stimulated cortisol levels are required to validate AI, with baseline cortisol levels serving only as a screening modality (43).
Figure 7. SSTR5 downregulates CRHR1 signaling via miRNA-449c. (A) Heatmap representing expression of candidate miRNAs that might bind Crhr1 3′ UTR in (lane 1) SSTR5-KO AtT20 cells, (lane 2) SSTR5-overexpressing AtT20 cells, and (lane 3) HP5 and WT mouse pituitaries. Asterisks identify 7 candidate miRNAs that showed similar trends of change similar in lane 2 and lane 3 and an opposite trend in lane 1. Candidate miRNA expression levels were analyzed by qPCR, and levels of each miRNA were normalized to endogenous U6 small nuclear RNA. (B) Crhr1 mRNA level measured in WT AtT20 cells (2.0 × 10^5 cells/well, n = 3) 48 hours after transfection with 200 nM miRNA inhibitors identified in A or scramble negative control. Crhr1 mRNA normalized to Gapdh and divided by the scramble control level. Experiments were repeated in quadruplicate, and representative results are shown. Nucleotide sequence of miRNA inhibitors is given in Supplemental Table 4. (C) Representative CRHR1 protein expression and quantitative densitometry analysis of receptor normalized to β-tubulin in WT AtT20 cells (n = 4) treated with miR-449c inhibitor. (D) CRH-stimulated ACTH levels measured by ELISA in the culture medium of whole pituitary glands (8.0 × 10^4 cells/well, n = 4 at each CRH concentration) collected from WT (n = 4) and HP5 (n = 6) mice. ACTH level (pg/ml) normalized to cell number (per 1,000 cells). Cells were transfected with 200 nM miR-449c inhibitor or scramble for 24 hours and treated with CRH for 120 minutes, and culture medium was collected. Y axis represents fold change in ACTH from respective CRH-untreated cells (NT). Each value represents the average of triplicate experiments. (E) miRNA-449c expression in AtT20 cells transfected with plasmid carrying miRNA-449c (pmR-449c) or empty plasmid (pmR-empty) normalized to endogenous U6 small nuclear RNA. Relative miRNA expression levels were normalized to Gapdh. Results represent the combination of 3 experiments performed in quadruplicate. (F) mRNA levels of indicated genes in AtT20 cells overexpressing miRNA-449c (pmR-449c) compared with cells expressing empty vector (pmR-empty). Results represent the combination of 3 experiments performed in quadruplicate. (G) SSTR5-KO #3 AtT20 cells transfected with pmR-449c or pmR-empty compared with similarly transfected WT cells for 48 hours and treated with increasing doses of CRH for 30 minutes, after which intracellular cAMP was measured by LANCE cAMP assay (8.0 × 10^4 cells/well, n = 4 for each CRH concentration). Y axis represents fold change in cAMP from control CRH-untreated cells (NT). Experiments were performed in triplicate. Results are presented as mean ± SEM. *P ≤ 0.05; **P ≤ 0.01; ***P ≤ 0.001; 2-tailed unpaired t test, (D and G) with Bonferroni correction.
Moreover, HP5 mice exhibit increased anxiety-like behavior, as demonstrated by less center activity and rearing behavior. Anxious behavior, a recognized feature of secondary AI (9, 50), is known to result from altered CRH signaling. Mice treated with a CRHR1 antagonist, as well as CRHR1-KO mice, display decreased anxiety (16, 51), while CRH-overexpressing Tg mice exhibit increased anxiety compared with CRH-KO mice (51, 52). The higher anxiety levels seen in HP5 mice likely results from a lack of negative feedback regulation on CRH, driven by reduced pituitary ACTH and peripheral corticosterone. Depressive mood, another behavioral symptom exhibited by patients with secondary AI (50), is also manifested in HP5 mice, which stopped swimming earlier on the forced swim test compared with WT mice. HP5 mice also exhibited greater deterioration in their health status upon treatment with LPS. HP5 mice are therefore a model for mild secondary AI, which present as clinically “normal” at baseline but show anxiety- and depressive-like behaviors and deteriorated health status upon exposure to an inflammatory challenge. The HP5 mouse is likely more clinically useful than other mouse models for simulating features of mild human secondary AI.

Several studies, including those from our laboratory, show SSTR5 activation leading to suppressed CRH signaling (29–31). In AtT20 cells that express SSTR2, SSTR3, and SSTR5, the SSTR1/SSTR5 selective agonist L-817,818 inhibited CRH-induced ACTH secretion (53). Similarly, pasireotide inhibited CRH-dependent ACTH secretion in rat, murine, and human pituitary cultures (30, 31). Constitutive SSTR5 activity in AtT20 cells also markedly suppressed CRH-induced ERK phosphorylation, as well as induction of intracellular cAMP levels and ACTH secretion in AtT20 cells (29).

Somatostatin directly inhibits exocytosis via inhibition of cAMP accumulation and calcium fluxes, thereby inhibiting hormone secretion (54). We now extend this understanding of somatostatin function by showing that SSTR5 negatively regulates ACTH production via posttranscriptional microRNA regulation of CRHR1. miR-34a is known to bind the 3’UTR of CRHR1 and, when overexpressed in human lymphocytes, downregulates CRHR1 expression and CRH signaling (55). In addition, miR-449a levels are increased in pituitary cells shortly after mice undergo a restraint test and/or are treated with dexamethasone, while overexpressed miR-449a suppresses CRHR1 (56). We found that, in mice overexpressing SSTR5, miR-449c specifically targets the CRHR1 3’UTR region. Consistent with these findings, HP5 mouse pituitaries and SSTR5-overexpressing AtT20 cells express higher levels of miR-449c and low CRHR1. The functional link between miR-449c and CRHR1 was further validated by showing that overexpressed miR-449c decreases CRHR1 expression, while miR-449c KO increases CRHR1 expression. Thus, we conclude that increased corticotroph SSTR5 and miR-449c expression act to downregulate CRHR1 posttranscriptional expression, reducing CRHR1 levels and corticotroph responses to CRH. Using this model, we elucidate a potentially novel mechanism by which SSTR5 controls CRH action and causes secondary AI.

It is possible that SSTR5 suppresses CRHR1 function by modifying the CRHR1 isoform expression profile, as several exhibit reduced activation of CRHR1 by CRH (57). Future studies will seek to define which CRHR1 isoform is involved in SSTR5 signaling through CRH.

A second potential limitation of our study is our reliance on 1 Tg founder. To minimize the possibility that expression profiles and responses were founder specific, we initially generated 3 separate Tg mice. All 3 showed similar expression of HA-hSSTR5 on IHC, and we selected 1 at random to serve as founder for further propagation.

In conclusion, we have developed a Tg mouse that demonstrates SSTR5 regulation of the HPA axis to generate biochemical and clinical features of secondary AI. We also show that, in addition to acute inhibition of pituitary ACTH secretion, SSTR5 posttranscriptionally suppresses CRHR1 via miR-449c, resulting in attenuated corticotroph responses to hypothalamic CRH. The HP5 mouse provides new insight into somatostatin receptor regulation of the HPA axis and recapitulates clinical and biochemical features of a mild form of secondary AI.

Methods

Study design
The objective of this study was to determine whether corticotroph SSTR5 attenuates the HPA axis stress response and to elucidate mechanisms by which it occurs. We evaluated the HPA axis stress response in a potentially novel corticotroph-specific SSTR5-overexpressing HP5 mouse model with different stress challenges, including biochemical stimulation (CRH), behavioral stimulation (restraint, open field test,
and forced swimming tests), and inflammatory challenge (LPS injection). To study mechanisms by which these occur, we assessed CRHR1 expression and response to SSTR5 levels in vivo, ex vivo, and in vitro (overexpression or KO of SSTR5 in AtT20 cells). As no effect on CRHR1 promoter activity was evident, we analyzed CRHR1-targeting miRNAs by qPCR, Western blotting, ELISA, cAMP assay, IHC, luciferase assay, and ChIP assay. Sample sizes for the animal studies were determined by the Biostatistics & Bioinformatics Core at Cedars-Sinai Medical Center.

**Treatments**

CRH (MilliporeSigma) and AVP (Abcam) were resuspended in sterile ddH₂O. LPS from *Escherichia coli* O111:B4 (MilliporeSigma) was resuspended in PBS. Forskolin (MilliporeSigma) was resuspended in dimethylsulfoxide. Pasireotide and octreotide (Novartis Pharmaceuticals) were resuspended in sterile ddH₂O. BIM-23206 (Ipsen) was prepared in 0.01 M acetic acid and 0.1% BSA as described (58).

**Generation of Tg (HP5) mice.** hSSTR5 cDNA was inserted downstream of the *Pomc*-enhancer-neuroD1-Tpit/Ptx1 (PENT) promoter in mouse pituitary corticotroph cells as described (36). cDNAs encoding hSSTR5 were provided by Graeme Bell (University of Chicago, Chicago, Illinois, USA). Xho I (5′ end) and EcoRV (3′ end) were introduced by PCR, and the hSSTR5 fragment joined upstream of HA ×3 sequences. The PENT promoter fragment in pGL4 vector was digested with NheI and XhoI and was inserted into the HA ×3 sequence containing hSSTR5 cDNA vector. PENT-hSSTR5 expression constructs were microinjected into male pronuclei of fertilized eggs, and injected eggs were transplanted to pseudopregnant foster mothers (FVB/NJ background; The Jackson Laboratory). Genomic DNA isolated from offspring 10–14 days old using a KAPA Mouse Genotyping Kit (KAPABIOSYSTEM) was analyzed by PCR using primers binding the POMC promoter (POMC-F: 5′ - GGTCACGTCCAAGGCTCACCCA - 3′) and SSTR5 cDNA (SSTR5-R: 5′ - CGTCCCTGTGTGTCACCCTCCA - 3′).

Backcross generations used in the present study ranged from F4–F9. Mice (females, 6–12 months old) were maintained in a pathogen-free temperature-controlled (23°C) mouse facility on the light/dark cycle (lights were on from 6 a.m. to 8 p.m.). Food and water were given ad libitum. Mice were fed with a normal diet (PicoLab Rodent Diet 20, Lab Supply). No more than 5 mice were housed per cage.

**Blood collection and hormone analyses.** Blood samples were collected by retro-orbital bleeding under anesthesia. After baseline samples were collected between 8 and 9 a.m. after an overnight fast, female mice were returned to their cages for 2 hours under fasting. Mice were injected i.p. with 40 μg/kg CRH. At 30 minutes after injection, blood samples were collected under anesthesia to obtain peak hormonal values.

**Restraint test.** On the first day, baseline blood samples were collected, female mice were returned to cages, and feeding resumed on a regular diet. On the second day, mice were manually restrained and placed into flat-bottomed restrainers (5 × 11 × 4 cm; Braintree Scientific Inc.) for 30 minutes without anesthesia. Immediately after removal from restraints, mice were euthanized, and blood and tissue were collected.

**LPS challenge test.** After baseline blood samples were collected, female mice were returned to cages for 2 hours under fasting and were then injected i.p. with LPS 5 mg/kg or 10 mg/kg. After 3 hours, retro-orbital blood was drawn from the opposite eye under anesthesia. Mice were then returned to cages, and feeding resumed. All mice were euthanized on day 3 after LPS administration, and blood and tissue were collected.

**Pasireotide injection test.** Experimental design is shown in Supplemental Figure 3A. Female mice were injected s.c. with pasireotide LAR 40 mg/kg under anesthesia at 9 a.m. every 2 weeks for a total of 3 injections (days 0, 14, 28). The CRH stimulation test was performed 1 week after the last injection.

**Open field test.** Female mice were moved into the test room in the Cedars-Sinai Rodent Bio-Behavior Core to habituate the day prior to the test. As this test measures spontaneous motor behavior, it was conducted first to avoid lingering effects from prior handling and manipulations. The open field apparatus is a 16-inch square, clear Plexiglas chamber with 15-inch high walls and an open top, surrounded by 2 rings of photobeams and optical sensors. The morning after transfer to the test room, mice were individually placed into the chamber, and photobeam breaks automatically recorded and were used to measure horizontal locomotor and rearing activity. Parameters recorded included general motor activity (number of beam breaks), number of times rearing, speed, distance traveled, and total resting time (59, 60).

**Forced swim test.** Female mice were moved into the test room in the Cedars-Sinai Rodent Bio-Behavior Core to habituate the day before the forced swim test, which measures behaviors that are indicators of despair in rodents (61, 62). The next morning, mice were individually placed into a large fully transparent container (40 cm height, 20 cm diameter) filled with water sufficient for the rodent to be fully submerged yet...
not to escape from the sides. Water temperature was maintained at 22°C–24°C, and mice were tested for 6 minutes. Mice were rated for immobility (defined as the absence of activity) and escape-oriented behaviors including swimming, climbing, and diving. Data were recorded and analyzed for 6 minutes.

Food consumption. Individual 3-month-old female mice were kept in metabolic cages (Nalgene Metabolic Cage - Single Mouse, Ancare Corp.) for 3 days for 24 hours to allow for acclimation and were then fed with Nutra-Gel Diet (Bio-Serv) to reduce the risk of sample contamination. Food consumption was measured by the difference in food weight before and after feeding for 2 days.

Cell cultures

AtT20. AtT20/D16v-F2 cells stably expressing hSSTR5-IRES-ZsGreen and AtT20/D16v-F2 cells expressing only pZsGreen1-N1 (29) were grown in serum-containing low-glucose DMEM (Invitrogen) supplemented with 10% FBS (Omega Scientifics), with the addition of 1% glutamine, 1% penicillin/streptomycin (Invitrogen) and genetin 400 μg/ml (Invitrogen), in 6% CO2 in a 37°C humidified incubator (29).

Primary mouse pituitary cell cultures. Primary mouse pituitary cells were isolated using the Neural Tissue Dissociation Kit (Miltenyi Biotec Inc.), plated in 48-well plates precoated with ECL Cell Attachment Matrix (MilliporeSigma), and attached after overnight incubation. Cells were cultured in serum-containing high-glucose DMEM (Invitrogen) supplemented with 10% FBS (Omega Scientifics), with the addition of 1% glutamine and 1% penicillin/streptomycin (Invitrogen), in 6% CO2 in a 37°C humidified incubator. Cells were then stimulated for 2 hours with increasing doses of CRH in serum-free medium, after which culture medium was collected for analysis. ACTH concentrations were corrected for cell number in each well.

KO of SSTR5 by CRISPR/Cas9

CRISPR/Cas9 reagents pSpCas9(BB)-2A-GFP (pX458, Addgene, 48138) were obtained from the Zhang laboratory via Addgene. Online CRISPR design tools were provided by Zhang Laboratory (http://crispr.mit.edu/) and CHOPCHOP (https://chopchop.rc.fas.harvard.edu/). Guides were selected for the fewest potential off-target sites and, where possible, the fewest potential off-target sites within coding exons. Guide sequences (sequence-PAM) were as follows: SSTR5KO#1 (exon 1, 40-bp deletion): A, GCACAGGCACTAATACCGCGG and B, GTTGGTAGGCACCGTTGGAC-TGGG; SSTR5KO#2 (exon 1, 37-bp deletion): A, CACAGGCACTAATACCGCGG and B, TACTTGTGGTGATGCAACCTTGGG; SSTR5KO#3 (exon 1, 47-bp deletion): A, AATACCGCCCGGGCTCCCAT-GGG and B, TACTTGTGGTGATGCAACCTTGGG. Guide sequences (without PAM) were cloned into pX458 and cotransfected into AtT20 cells using Neon Transfection System (Invitrogen) using 1150 V for 30 ms in 2 pulses. After 48 hours, transfected cells were individually placed into single wells of a 96-well plate. Isolated cell clones were grown and tested using genotyping primers (forward, 5'- GCTTCCAGCAGTAGCCATAA - 3' and reverse, 5'- CAGGAAAGGAAGCCCCAACA - 3') that spanned the deleted genomic region. Clones were also PCR amplified and sequenced to verify the out-of-frame deletion in genomic material. The SSTR5 mutant clones were confirmed with disruption of SSTR5 function on intracellular cAMP assay using forskolin and BIM-23260 (Supplemental Figure 4E).

Transfections

Transfections of plasmid or microRNA inhibitors to cells were performed by Neon Transfection System (Thermo Fisher Scientific) at room temperature using 1150 V for 30 ms in 2 pulses for AtT20 cells or 1700 V for 20 ms in 1 pulse for primary mouse pituitary cells, with 100 μl R buffer for 2.0 × 10^6 cells/each reaction.

Immunoblotting

Total cell lysate was prepared in RIPA buffer (MilliporeSigma) containing Protease Inhibitor Cocktail (MilliporeSigma). For Western blot analysis, proteins were separated by SDS/PAGE, electroblotted onto TransBlot Turbo Transfer Pack 0.2-μm PVDF membrane (Bio-Rad), and incubated with Blocking One (Nacalai USA Inc.) for 20 minutes; they were then incubated overnight with primary antibodies at 4°C. The following day, membranes were incubated with corresponding secondary antibodies for 2 hours at room temperature. Both primary and secondary antibodies were resuspended into Signal Enhancer HIKARI (Nacalai USA). Protein concentration was measured by DC protein assay kits (Bio-Rad), and absorbance was measured at 750 nM with the Victor^2 spectrophotometer (Perkin Elmer). Loading quality was confirmed by Ponceau S (MilliporeSigma) staining.
Immunofluorescence

Mouse pituitary glands were fixed in 10% formalin and embedded in paraffin. After deparaffinization and antigen retrieval, slides were blocked in Blocking One Histo (Nacalai USA), incubated overnight with rabbit polyclonal anti-hSSTR5 antibodies (1:100, Abcam; catalogs ab109495 and ab138863, 1:100, Santa Cruz Biotechnology Inc.; catalog sc-25679) and mouse monoclonal anti-ACTH (1:500, Abcam; ab20358). After washing, samples were incubated with Alexa Fluor chicken anti–mouse 488 (H+L, 1:500 dilution, Invitrogen; catalog A-21200) and Alexa Fluor chicken anti–mouse 647 (H+L, 1:500 dilution, Invitrogen; catalog A-21463) mounted with Prolong Gold antifade reagent (Invitrogen). Confocal microscope images were obtained using True Confocal Scanner (Leica Microsystems) in dual-emission mode to separate auto-fluorescence from specific staining.

Antibodies

Anti-CRHR1 antibody (1:1,000, Aviva Systems Biology; catalog OASG01837), anti-hSSTR5 (1:1,000, Abcam; catalog ab109495) (63, 64), HA-tag (1:1,000, Cell Signaling; catalog 3724), β-tubulin (1:1,000, Abcam; catalog ab6046), rat GH (1:3,000, R&D Systems; catalog MAB1566), ACTH (1:2000, Abcam; catalog ab20358), PC1/3 (1:1,000, Cell Signaling Technologies; catalog 11914S), PC2 (1:1,000, Cell Signaling Technologies; catalog 14013), AVP (1:1,000, MilliporeSigma; catalog AB1565), HA (1:1,000, Cell Signaling Technologies; catalog 3724), and c-Myc (1:1,000, Cell Signaling Technologies; catalog 5605) were purchased. Tpit antibody (1:1,000) was provided by Jacques Drouin (Clinical Research Institute of Montreal, Montreal, Quebec, Canada).

Assays

Hormones were assessed using ACTH ELISA kit (MD Biosciences Inc.), Corticosterone Double Antibody RIA Kit (MP Biomedicals), αMSH (Human, Rat, Mouse) EIA Kit (Phoenix Pharmaceuticals Inc.), Mouse/Rat IGF-I ELISA Kit (ALPCO Ltd.), Prolactin Mouse ELISA kit (Thermo Fisher Scientific), Rat/Mouse Insulin ELISA (LINCO Research), and One Touch Ultra for glucose (Johnson & Johnson).

**cAMP assay.** Cells seeded overnight at a density of 8 × 10^4 cells/well in a 48-well plate were stimulated for 30 minutes at 37°C in medium containing 0.3% BSA and 1 mM 3-isobutyl-1-methylxanthine (MilliporeSigma) supplemented with indicated treatments. Duplicate samples were assayed for cAMP levels with the LANCE cAMP Kit (PerkinElmer) and modified for intracellular cAMP measurement. Results were extrapolated from standard curves measured by the Victor^2 spectrophotometer (PerkinElmer) (65).

**Luciferase assay.** The mouse Crhr1 promoter (−1,400 to +68) was PCR amplified from genomic DNA (AtT20 cells) and cloned. Promoter regions were amplified using 5′ primers binding to −1,334, −912, −728, −466, and −266 (containing Xho I site at the 5′ end) and 3′ primer binding to −13 (containing Hind III sites at the 3′ end). Resulting amplified fragments were digested with Xho I and Hind III and cloned into Xho I and Hind III sites in pGL3 basic vector. DNA sequences of inserted fragments were determined to remove defective fragments generated by PCR errors. Luciferase reporter assays were performed in 8 × 10^4 AtT20 cells with 0.5 μg luciferase reporter plasmids, and 10 ng pRL-CMV was used as an internal control plasmid. Cells were transfected using lipofectamine 2000 (Invitrogen) and cultured in 1 ml medium in 48-well plates. Forty-eight hours after transfection, cells were harvested, and luciferase activity was analyzed by Dual-Luciferase Reporter Assay System (Promega). Assays were repeated at least 3 times, and luciferase activity was normalized by internal renilla activity.

**ChIP assay.** Two million AtT20 cells were fixed for 10 minutes at room temperature in 1% formaldehyde, 4.5 mM HEPES (pH 8.0), 9 mM NaCl, 0.09 mM EDTA, and 0.045 mM EGTA and sonicated (Nanoruptor, NR-350, Cosmobio Co. Ltd.) in lysis buffer (1% SDS, 10 mM EDTA, and 50 mM Tris–HCl (pH 8.0)) with proteinase inhibitor (MilliporeSigma P8340). ChIP Assay Kit (MilliporeSigma) was used, and precleared lysates were incubated overnight at 4°C with polyclonal anti-acetyl histone H4 (Millipore) or control rabbit IgG (Santa Cruz Biotechnology Inc.). DNA fragments were isolated from immunoprecipitated chromatin and analyzed by real-time PCR with SsoAdvance SYBR Supermix (Bio-Rad). ChIP PCR primers are shown in Supplemental Table 1.

RNA extraction and real-time PCR

RNA was collected with RNeasy Mini Kit (QIAGEN). cDNA was synthesized from 1 μg purified RNA by the iScript Reverse Transcription Supermix for qPCR (Bio-Rad). qPCR was performed in 10 μl reactions...
using SsoAdvanced Universal SYBR Green Supermix in CFX96 or 384 instrument (Bio-Rad). Nucleotide sequence of primers is given in Supplemental Table 2. TaqMan Gene Expression Assays (Applied Biosystems) were used to analyze hSSTR5 expression (Hs00990408_s1). TaqMan mouse GAPDH Expression Assays with probe tagged with VIC (Applied Biosystems) were used as reference genes.

MicroRNA

Analysis. Total RNA for miRNA expression analysis was extracted from cells using TRIzol reagent (Invitrogen). Mir-X miRNA First-Strand Synthesis Kit and Mir-X miRNA qRT-PCR SYBR Kit (Clontech; Takara Bio Inc.) were used to analyze miRNA expression. miRNA primer sequences were obtained from miRBase (http://www.mirbase.org/) (Supplemental Table 3). Ct values >35 were removed from analysis. Expression levels were normalized by U6 with primers provided in the Mir-X miRNA qPCR SYBR Kit (Clontech; Takara Bio Inc.).

Overexpression and inhibition of miRNA. mmu-miR-449c (NR_030452.1) was PCR amplified using a set of primers (forward, 5′- GCTAGATCTAGGATGAAGTGTGGGTGTG - 3′, containing Bgl II site at the 5′-end, and reverse, 5′- CACGGATCCGCATGCTTCTGAGTG - 3′, containing BamH I sites at the 3′-end) from genomic mouse DNA and cloned. Resulting amplified fragments were digested with and cloned into Bgl II and BamH I in pmR-ZsGreen1 miRNA overexpression vector (Clontech; Takara Bio Inc.). Steric blocking miRNA inhibitors that hybridize to mature miRNAs and NC5 negative control were purchased from Integrated DNA Technologies Inc. Primary mouse pituitary cells or AtT20 cells were transfected with our constructs by Neon Transfection System (Invitrogen) using 1150 V for 30 ms in 2 pulses for AtT20 cells and 1700 V for 20 ms in 1 pulse for primary mouse pituitary cells, both with 100 μl R buffer. Experiments were performed after 48 hours.

Statistics

Statistical analyses were performed with GraphPad Prism 7.0 (GraphPad Software Inc.). Comparisons were analyzed by unpaired 2-tailed Student’s t test. Multiple-comparisons were analyzed by unpaired 2-tailed Student’s t test with the Bonferroni correction. P ≤ 0.05 was considered statistically significant.

Study approval

Animal care and handling was approved by the Cedars-Sinai IACUC (protocol no. 3392) and in accordance with the Guide for the Care and Use of Laboratory Animals (National Academies Press, 2011).

Author contributions

ABS and SM designed the study, and MY and HK performed the experiments. MY, ABS, HK, ND, and SM analyzed the data, and MY, ABS, and SM wrote the manuscript. HF created and provided plasmid used to express SSTR5 in corticotroph cells, and ABS generated the Tg mice. ED assisted with sample collection and provided technical assistance.

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