Satb2 regulates the development of dopaminergic neurons in the arcuate nucleus by Dlx1

Qiong Zhang1,2,3,7, Lei Zhang2,4,7, Ying Huang1, Pengcheng Ma5,6, Bingyu Mao5,6, Yu-Qiang Ding2,3 and Ning-Ning Song1,3✉

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INTRODUCTION

Hypothalamus, in particular the arcuate nucleus (ARC), is critical for the regulation of feeding and metabolism [1]. Two major types of neurons are identified in the ARC: neurons expressing the anorexigenic α-melanocyte-stimulating hormone derived from POMC, and those expressing AgRP or NPY [2]. Generally, POMC neurons promote activity, energy expenditure and suppress food intake, while AgRP/NPY neurons suppress energy use and promote food intake. In addition, the ARC also contains other types of neurons, such as those expressing somatostatin (Sst), growth hormone-releasing hormone (GHRH), or galanin in the ARC. Nestin-Cre;Satb2lox/fox (Satb2 CKO) mice show a reduced number of ARC DA neurons with unchanged numbers of the other types of ARC neurons, and exhibit an increase of serum prolactin level and an elevated metabolic rate. The reduction of ARC DA neurons in the CKO mice is observed at an embryonic stage and Dlx1 is identified as a potential downstream gene of Satb2 in regulating the development of ARC DA neurons. Together, our study demonstrates that Satb2 plays a critical role in the gene regulatory network directing the development of DA neurons in ARC.

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1Department of Laboratory Animal Science, Fudan University, Shanghai, China. 2Key Laboratory of Arrhythmias, Ministry of Education of China, East Hospital, and Department of Anatomy and Neurobiology, Tongji University School of Medicine, Shanghai, China. 3State Key Laboratory of Medical Neurobiology and MOE Frontiers Center for Brain Science, Institutes of Brain Science, Fudan University, Shanghai, China. 4Department of Anatomy, Histology and Embryology, School of Basic Medical Sciences, Fudan University, Shanghai, China. 5State Key Laboratory of Genetic Resources and Evolution, Kunming Institute of Zoology, Chinese Academy of Sciences, Kunming, China. 6Center for Excellence in Animal Evolution and Genetics, Chinese Academy of Sciences, Kunming, China. *These authors contributed equally: Qiong Zhang, Lei Zhang. ✉email: songnn1982@outlook.com

Dopaminergic (DA) neurons in the arcuate nucleus (ARC) of the hypothalamus play essential roles in the secretion of prolactin and the regulation of energy homeostasis. However, the gene regulatory network responsible for the development of the DA neurons remains poorly understood. Here we report that the transcription factor special AT-rich binding protein 2 (Satb2) is required for the development of ARC DA neurons. Satb2 is expressed in a large proportion of DA neurons without colocalization with proopiomelanocortin (POMC), orexigenic agouti-related peptide (AgRP), neuropeptide-Y (NPY), somatostatin (Sst), growth hormone-releasing hormone (GHRH), or galanin in the ARC. Nestin-Cre;Satb2lox/fox (Satb2 CKO) mice show a reduced number of ARC DA neurons with unchanged numbers of the other types of ARC neurons, and exhibit an increase of serum prolactin level and an elevated metabolic rate. The reduction of ARC DA neurons in the CKO mice is observed at an embryonic stage and Dlx1 is identified as a potential downstream gene of Satb2 in regulating the development of ARC DA neurons. Together, our study demonstrates that Satb2 plays a critical role in the gene regulatory network directing the development of DA neurons in ARC.

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revealed that Dlx1 is likely to be a downstream gene of Satb2 in regulating the development of ARC DA neurons.

RESULTS
Lower body mass and higher metabolic rate in Satb2 CKO mice
Previous studies have shown that Satb2 is critical for cortical development [11–14, 16–18]. However, the role of Satb2 in other brain regions is largely unknown. To this end, Satb2 floxed mice [19] were crossed with Nestin-Cre mice [20] to knock out this gene in the central nervous system. Littermates with other genotypes (i.e., Satb2flox/+ and Satb2flox/flox) had no alterations examined below and were used as controls. Satb2 CKO mice were viable and could survive into the adult stage, but showed reduced body weight during postnatal development and at adult and aged stages compared to their littermate controls (Fig. 1a). To investigate if the lower body weight was due to changes in basal metabolic rate, we placed the Satb2 CKO and control mice in the metabolic chambers and measured the oxygen consumption (VO2) and carbon dioxide expiration (VCO2) for 24 h by indirect calorimetry. Consistent with the reduction in body weight, Satb2 CKO mice exhibited a significant increase in VO2 during the total 24 h period including the dark and light cycles (Fig. 1b), indicating a higher basal metabolic rate in these mice. However, we did not detect a change in the respiratory exchange rate (RER, VCO2/VO2) (Fig. 1c) or energy expenditure (Fig. 1d). Taken together, our data revealed a higher metabolic rate and growth retardation in Satb2 CKO mice.

DA neurons are selectively reduced in ARC of adult Satb2 CKO mice
Several hypothalamic nuclei including the ARC sense nutrient and endocrine cues and coordinate metabolic responses [21]. Our previous study showed that Satb2 is expressed in the ARC in adult mice [15]. Thus, the dysregulation of metabolism in Satb2 CKO mice prompted us to examine if there are any changes of the neuron types involved in metabolic activity, including POMC, NPY/AgRP, and tyrosine hydroxylase (TH, a marker for DA neurons) positive neurons. Using immunostaining, we first confirmed the expression of Satb2 in the ARC of control mice and verified the deletion of this gene in Satb2 CKO mice at P60 (Fig. 2a, a’). The Nissl-stained cellular architecture of the ARC was well maintained in Satb2 CKO mice relative to controls (Fig. 2b, b’). Then, we determined the neuron types by in situ hybridization (ISH). The quantitation results showed that the numbers of POMC+ (Fig. 2c, c’, i) and NPY+ (Fig. 2d, d’, j) neurons were comparable in the ARC between control and Satb2 CKO mice, but the ISH signals of NPY with the same duration for signal development (1 h) was stronger in the ARC of Satb2 CKO mice than that in control (Fig. 2d, d”, k). Similar results were obtained for the ISH of AgRP in ARC (Supplementary Fig. S1). The number of TH+ neurons was significantly reduced in both DM and VL parts of ARC in Satb2 CKO mice, although the reduction in the VL was less severe than DM (Fig. 2e, e’, l). To exclude the possibility that the reduction is caused by the inability of TH expression itself rather than decreased neuron number, we examined the expression of other DA neuron markers including vesicular monoamine transporter 2 (VMAT2), aromatic L-amino acid decarboxylase (AADC), and

![Fig. 1](https://example.com/fig1.png)  
**Fig. 1**  
Lower body mass and higher metabolic rate in Satb2 CKO mice.  
a The body weight of Satb2 CKO mice is sharply reduced when compared with control mice during postnatal development, at adulthood, and at the aged stage. At P7-P60, \( n = 8 \) for control and \( n = 6 \) for Satb2 CKO mice; at P300, \( n = 11 \) for control and \( n = 7 \) for Satb2 CKO mice. Error bars represent S.D.  
b Oxygen consumption (VO2) was measured by indirect calorimetry in control and Satb2 CKO mice. Error bars represent S.E.M. Statistical differences were determined by Student’s t-test, *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001; n.s. not statistically significant.
dopamine transporter (DAT). Similar to the changes in TH expression, the expression of VMAT2, AADC and DAT were also dramatically decreased in the ARC of Satb2 CKO mice (Fig. 2f–h'). Thus, the number of DA neurons is indeed reduced with no obvious changes in the numbers of NPY⁺ and POMC⁺ neurons in the ARC of Satb2 CKO mice.

Satb2 is expressed in adult and developing ARC DA neurons

To explore why only TH expression was reduced in the ARC, we checked the colocalization of Satb2 with these genes in wild-type mice. Immunostaining of Satb2 combined with ISH of AgRP or NPY showed no colocalization of Satb2 with these peptides (Fig. 3a, b). Meanwhile, Satb2 was not detected in POMC⁺ neurons in the ARC of Satb2 CKO mice.
POMC-GFP mice [22] (Fig. 3c). Consistent with our previous data [15], TH/Satb2 double-labeled neurons were observed in the ARC at P60 (Fig. 3d–f). We quantified the proportion of Satb2/TH-coexpressing neurons, and about 88% of TH+ neurons expressed Satb2, which corresponded to about 57% of Satb2+ neurons in the DM. In the VL, about 69% of TH+ neurons expressed Satb2, which was present in about 59% of Satb2+ cells (Fig. 3n). These data indicate that Satb2 is expressed in ARC DA neurons in both DM and VL with different proportions. Taken together, these data suggest an autonomous role of Satb2 in regulating the development of ARC DA neurons.

We next started to explore the expression pattern of Satb2 in developing ARC DA neurons. We investigated its expression in the ARC at different embryonic stages in combination with two specific markers Cbln1 [23] and POMC [24], which label the ventromedial hypothalamic nucleus (VMH) and ARC, respectively. Satb2 was first observed in the presumptive hypothalamus at E12.5 (Fig. 3e, f), and the vast majority of Satb2+ cells were located dorsally to POMC+ ARC domain (Fig. 3e) and within the Cbln1+ VMH domain (Fig. 3f). At E14.5, in addition to the presence of numerous Satb2+ cells in the VMH, a few Satb2+ cells were found to be located ventrally to the Cbln1+ domain and within the ARC region containing POMC+ cells (Fig. 3g, h). To determine Satb2 is expressed in progenitors or postmitotic cells, BrdU was injected intrauterinely 1 h before the sacrifice of the pregnant mice. No Satb2+ cells were labeled with BrdU at E12.5, E14.5, and E16.5 (Fig. 3i, j, and data not shown), indicating that Satb2 is expressed in postmitotic ARC neurons.

We next investigated the expression of Satb2 in comparison with that of TH at different embryonic stages. Immunostaining of Satb2 was performed in adjacent sections of those processed for detection of TH mRNA at E14.5, 15.5, and 16.5. TH expression was initiated in the VL at E14.5 and DM at E15.5 in the ARC (Fig. 3k–m). However, Satb2 expression was first detected in the ARC at E14.5, and it was located in the DM only (Fig. 3k’, o). Satb2 was first detected in the VL at E15.5 and its expression was increased in both DM and VL at E16.5 (Fig. 3l, m’, o). Colocalization of Satb2 and TH was observed at E16.5 and P0 (Fig. 3p–s). This unique expression pattern continued to P0 (Fig. 3s) and P60 (Fig. 3d). Thus, the initiation of Satb2 expression in the DM occurs earlier (i.e., E14.5) than that of TH (i.e., E15.5), but those in the VL show the opposite way during embryonic development.

Satb2 is required for early differentiation of DA neurons in DM and for maintenance of DA neurons in VL

The decline of DA neurons in adult Satb2 CKO mice might be due to defective generation and/or impaired maintenance. To figure out which was the case, we examined TH expression at different embryonic stages in Satb2 CKO mice. TH transcript was detected in the VL of control mice at E14.5 (Fig. 4a), and a similar expression pattern was observed in Satb2 CKO embryos (arrows, Fig. 4a, a’), showing that the initiation of TH expression in the VL is not affected. At E15.5, TH expression in the VL was similar between control and Satb2 CKO mice, but a significant reduction was observed in the DM of Satb2 CKO mice (Fig. 4b, b’), suggesting that the initiation of TH expression in the DM is impaired. At E16.5, an apparent increase of TH transcripts was observed in the DM of control mice but not in CKO mice, while the expression of TH mRNA seemed to be unchanged in the VL of Satb2 CKO mice relative to controls (Fig. 4c, c’). An obvious reduction of TH expression in the VL was found in Satb2 CKO mice at P14, and few TH-expressing cells were observed in the DM at this stage (Fig. 4d, d’). Thus, defective DA neuron differentiation occurs in the DM as early as E14.5, while a failure of DA neuron maintenance starts in the VL at postnatal stages in Satb2 CKO mice.

GAD67+, Sst+, GHRH+, and Galanin+ neurons remain unchanged in ARC of Satb2 CKO mice

We have shown that the development of DA neurons is affected in Satb2 CKO mice with unchanged NPY− and POMC+ populations. However, there are several other types of neurons in the ARC characterized by distinct neurotransmitters, neuroendocrine functions, or expression of specific neuropeptides, including GAD67, Sst, GHRH, and Galanin [25, 26]. We next examined if these neuronal populations were altered by double immunostaining or ISH combined with immunostaining in wild-type mice. The majority of Satb2+ neurons expressed GFP in the ARC of GAD67-GFP mice [27] (Fig. 5a, i). Similar to the cases of AgRP/NPY/POMC, there were no Satb2+ neurons expressing Sst, GHRH, or Galanin (Fig. 5b–d). The number of GAD67+ cells in the ARC of Satb2 CKO mice was not different from that of control mice (Fig. 5e, e’, j). As expected, cell counts showed that the numbers of Sst+, GHRH+, and Galanin+ neurons were not significantly changed in adult Satb2 CKO mice compared with controls (Fig. 5f–h’, k–m). Taken together, our data indicate that Satb2 is selectively involved in the development of ARC DA neurons.

As mentioned above, Satb2 is also expressed in the VMH (Fig. 3). However, the cellular architecture of VMH shown by Nissl staining was not apparently altered in adult Satb2 CKO mice (Fig. 2b, b’). Furthermore, several genes are specifically expressed in the VMH such as Cbln1, Nr5a1, and Sox14 [23, 26], and their expression was also comparable between control and Satb2 CKO mice (Supplementary Fig. S2), suggesting that the overall morphology of the VMH might not be significantly affected. In addition, cleaved Caspase3 immunostaining at E14.5, E16.5, and P0 did not detect significant differences of the positive cells in the ARC between control and Satb2 CKO mice (Supplementary Fig. S3).

Serum prolactin level is increased in female Satb2 CKO mice

ARC DA neurons serve as an inhibitor in regulating the secretion of prolactin [29]. We thus set out to examine the expression of pSTAT5, a signal transducer and activator for prolactin secretion [30] in virgin and lactating females. Immunostaining showed that the expression of pSTAT5 was dramatically increased in the ARC and VMH in lactating dams compared with virgin females (Fig. 6a, a’), consistent with the notion that suckling stimulates the
secretion of prolactin in lactation [29]. The proportion of pSTAT5 expression in DA neurons was also significantly increased in the DM and VL of ARC in lactating mice (inserts in Fig. 6c, c', d). Consistent with the inhibitory role of ARC DA neurons in regulating the secretion of prolactin, we found that the levels of serum prolactin were increased in adult female Satb2 CKO mice, although those in male CKO mice showed an increasing trend (Fig. 6e). Besides, pSTAT5 was abundantly expressed in the ARC, VMH...
Satb2 is required for the differentiation of DA neurons in DM and for the maintenance of DA neurons in VL. a–c’ Comparison of TH expression in the ARC of control and Satb2 CKO mice at different embryonic stages. TH mRNA is detected in the VL (arrows) in both control (a–c) and Satb2 CKO (a’–c’) embryos with no obvious differences at E14.5, E15.5, or E16.5. TH mRNA is present in the DM (triangles) of control mice at E15.5 (b) and E16.5 (c) whereas no TH transcripts are found in age-matched Satb2 CKO mice (b’, c’). d, d’ Reduction of TH expression is first detected in the VL (arrow) of Satb2 CKO mice at P14, when few TH+ neurons are present in the DM (triangle) relative to controls. 3V third ventricle. Scare bars = 100 μm in a–d’.

and medial nucleus of the amygdala (MEA) of female Satb2 CKO mice (inserts in Fig. 6f, g’), whereas few pSTAT5+ cells were scattered in these regions in control mice (inserts in Fig. 6f, g), which is consistent with the increase of prolactin secretion in Satb2 CKO mice. In addition, the size of the pituitary gland was reduced in Satb2 CKO mice compared with control mice (Supplementary Fig. S4a–c), which is consistent with the reduction of body weight in Satb2 CKO mice (Fig. 1a). The secretion of prolactin is regulated by ARC-dopamine through the D2 receptor (DRD2), which is expressed in both the intermediate lobe (IL) and anterior lobe (AL) of the pituitary gland [31]. ISH showed that the expression of Drd2 in the IL and AL of the pituitary gland (31), but the intensity of GH immuno-fluorescence was not obviously changed in the AL of Satb2 CKO mice compared with control, suggesting that the generation of GH may not be affected (Supplementary Fig. S4d, g). Collectively, it is very likely that reduced ARC DA neurons lead to elevated prolactin levels in Satb2 CKO mice.

**Dlx1 acts downstream of Satb2 in development of ARC DA neurons**

Previous studies have identified several transcription factors that are involved in the development of the ARC [7–10, 32]. We found that Nkx2.1 expression in the ventral hypothalamus was not altered in the CKO mice relative to controls (Supplementary Fig. S5a, a’). In addition, the expression of Islet1 was not changed either in the ARC of Satb2 CKO mice (Fig. 7a, a’). Further, Mash1 expression was not changed in Satb2 CKO mice (Supplementary Fig. S5b, b’). We next investigated the expression of Dlx1 in the ARC. Dlx1 expression was not different in the ARC between control and Satb2 CKO mice at E14.5 (Fig. 7b, b’). However, its expression was dramatically reduced in the DM with a less severe reduction in the VL of Satb2 CKO mice at E16.5 (Fig. 7c, c’), which resembled the alternation of TH expression in the ARC. This change was confirmed by ISH for Dlx1 mRNA (Fig. 7d, d’). The quantitation of Dlx1+ cells in the DM and VL of ARC at E16.5 is shown in Fig. 7e. The loss of DA neurons in the ARC has been reported in Dlx1 KO and CKO mice [10, 33], and here we showed that Dlx1 expression depends on Satb2 in the ARC, raising the possibility that Satb2 regulates ARC DA neuron development by Dlx1.

To explore this, we examined the colocalization of Dlx1 and Satb2 in the ARC. The majority of Dlx1+ cells expressed Satb2 in the DM and to less extent in the VL at E16.5 (Fig. 7f–h’). In the DM, 93% of Dlx1+ cells expressed Satb2, while 86% of Satb2+ cells expressed Dlx1. However, in the VL, 33% of Dlx1+ cells expressed Satb2, while 90% of Satb2+ cells expressed Dlx1 (Fig. 7i). In addition, we also found that most TH+ cells expressed Dlx1 in both DM and VL of ARC at P7 (Supplementary Fig. S6a–c’). Satb2 is known to interact with nuclear matrix attachment regions (MAR) and regulate transcription of downstream genes via direct binding to MAR sites [34]. To determine whether Satb2 protein directly binds to MAR sites of the Dlx1 gene and regulates its expression, we performed ChIP and dual-luciferase reporter assays. According to theoretical prediction [http://genomecluster.secs.oakland.edu/marwiz/], there are three putative MAR sites in the Dlx1 genome (Fig. 7j). ChIP assay showed significant enrichment of these Dlx1-MAR sequences after Satb2 antibody incubation (Fig. 7k), indicating a direct association of Satb2 protein with Dlx1-MAR sequences. Then the dual-luciferase reporter assay was performed to examine the transcription activity of Satb2 in regulating Dlx1 expression. When Dlx1-MAR sequences were cloned into pGL4.73 plasmid, luciferase activity was increased when Satb2 cDNA was co-transfected (Fig. 7l). Taken together, these data suggest that Dlx1 may function as a downstream gene of Satb2 in regulating the development of DA neurons in the ARC.
DISCUSSION

The ARC has a critical role in the regulation of feeding and energy balance. Two distinct and functionally antagonistic types of neurons positive for POMC or AgRP/NPY are ‘first-order’ neurons in the ARC, which receive and integrate metabolic signals [35]. We did not detect any changes in the cell number of POMC or AgRP/NPY neurons in Satb2 CKO mice (Fig. 2). Recently, DA neurons that control the activity of adjacent POMC and AgRP/NPY neurons and thus are involved in the regulation of energy homeostasis have been identified in the ARC [6]. Consistently, we found that DA neurons are significantly decreased in Satb2 CKO mice (Figs. 2 and 4). Besides, the ARC DA neurons are well-known for a suppressor role in the regulation of the secretion of prolactin via the tuberoinfundibular pathway [29]. We found that the serum prolactin level is increased, and prolactin-responsive neurons shown by pSATA5 are also increased in the brain of Satb2 CKO mice, similar to the elevation of pSATA5 in lactating females (Fig. 6). Although the expression of GH is unchanged ([Supplementary Fig. 5]

**Fig. 5  GAD67, Sst, GHRH, and Galanin-positive cells remain unchanged in ARC of Satb2 CKO mice.** a Satb2 immunoreactivity is observed in a population of GFP⁺ cells in the ARC of adult GAD67-GFP transgenic mice. b–d Immunostaining of Satb2 combined with ISH of Sst, GHRH, and Galanin in wild-type mice. No colocalization of Satb2 with these peptides is observed. e–h’ Expression GAD67, Sst, GHRH, and Galanin in the ARC is comparable between adult control (e–h) and Satb2 CKO mice (e’–h’). i The quantification of co-labeling of Satb2 and GAD67-GFP in the ARC of adult GAD67-GFP transgenic mice. n = 2. j–m The quantification of GAD67⁺ (j), Sst⁺ (k), GHRH⁺ (l), and Galanin⁺ (m) cells in the ARC shows no significant differences between the two groups. n = 3 for each group. Statistical differences were determined by Student’s t-test. Error bars represent S.E.M. 3V third ventricle; ARC arcuate nucleus; VMH ventromedial hypothalamic nucleus. Scale bars = 100 μm in a, 100 μm in b–d, 25 μm in inserts of b–d, and 100 μm in e–h’.

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Fig. S4), the reduced size of the pituitary gland may reflect defective synthesis and secretion of other hormones in the gland. Thus, further studies are needed to explore if the other hormones are changed and their contributions to dysregulation of body weight and metabolism observed in Satb2 CKO mice.

Previous studies have identified several proneural genes that are involved in controlling the development of ARC in the hypothalamus, such as Mash1 [7], Ngn3 [36], and Nkx2.1 [8]. Our results showed that Satb2 is only expressed in postmitotic neurons of ARC (Fig. 3), and expression of Nkx2.1 and Mash1 is not significantly changed in Satb2 CKO mice (Supplementary Fig. S5). Given that the apoptosis is comparable in the ARC between control and Satb2 CKO mice, it thus is likely that the reduction of DA neurons in the ARC is caused by defective differentiation of postmitotic DA neurons in the absence of Satb2. However, it should also be noted that the DA neurons in the ARC are not equally affected in Satb2 CKO mice, as shown by a more drastic reduction of DA neurons in the DM than that in the VL. A higher proportion of DA neurons in the DM express Satb2 in comparison with that in the VL (88% in DM versus 57% in VL), and this may account for the discrepancy, indicating that Satb2 is required for the differentiation of DA neurons in a cell-autonomous way. On the other hand, Satb2 seems to regulate the differentiation of ARC DA neurons in different ways. The initiation of Satb2 is prior to and required for that of TH in the DM. In the VL, however, Satb2 and TH initiate in an opposite way, and the initiation of TH is normal in Satb2 CKO mice. The reduction of DA neurons in the VL occurs during early postnatal days and this supports the idea that Satb2 is required for the maintenance of the neurotransmitter phenotype of DA neurons in the ARC.

In summary, we found that the body weight and metabolic rate are altered in Satb2 CKO mice, possibly due to the reduction of ARC DA neurons. We also showed that Satb2 regulates the development of ARC DA neurons through potential downstream gene Dlx1. We finally revealed one of the functional consequences of impaired ARC DA neurons shown by the increased level of serum prolactin.

**MATERIALS AND METHODS**

**Animals**

All mice were housed in a specific pathogen-free animal facility under a normal 12 h light, 12 h dark cycle with ad libitum access to normal chow and water. To conditionally knock out Satb2 gene in the brain, Nestin-Cre
mice [20] were crossed with floxed Satb2 mice [19] to delete exon 4 (Nestin-Cre;Satb2<sup>fl</sup>ox/fl<sup>ox</sup>, referred to as Satb2 CKO thereafter). POMC-GFP and GAD67-GFP mice were used as described previously [22, 37]. For virgin and lactating mice, sexually naïve females were used. The lactating group was mated with males and the virgin group was not. About 10 days after the birth of pups, the lactating mice were euthanized for experiments, along with the virgin mice.

Metabolic expenditure measurements
The control and Satb2 CKO mice of 10-month-old were placed in a Comprehensive Laboratory Animal Monitoring System (CLAMS, Columbus Instruments, Columbus, OH) for the evaluation of metabolism. Oxygen consumption (VO2) and carbon dioxide expiration (VCO2) were measured. The respiratory exchange ratio (RER) and energy expenditure were calculated based on the oxygen and carbon dioxide data.

Immunohistochemistry, BrdU labeling, and in situ hybridization
The stage of mouse embryos was determined by taking the morning when the copulation plug was observed as embryonic day 0.5 (E0.5). Anesthetized embryos and mice were perfused transcardially with phosphate-buffer saline (PBS; pH7.4) first then with 4% paraformaldehyde (PFA). All brains were fixed in 4% PFA overnight and cryoprotected in PBS containing 30% sucrose. Brains were cut into 20-μm-thick sections using a cryostat (CM1950; Leica, Wetzlar, Germany).

For immunohistochemistry, brain sections were then incubated with rabbit anti-Satb2 (1:300; ab92446, Abcam, Cambridge, UK), mouse anti-Satb2 (1:200; ab51502, Abcam), rat anti-BrdU (1:300; OBT0030, Accurate Chemical & Scientific Corporation, Carle Place, NY), Cleaved Caspase3 (1:300; 9661, Cell Signaling Technology, Danvers, MA), pSTAT5 (1:500; #9351, Cell Signaling Technology), Islet1 (1:100; 40.2D6, DSHB, Iowa City, USA), and Dlx1 antibodies (1:300; sc85510, Santa Cruz Biotechnology, Dallas, TX).

For in situ hybridization, brain sections were hybridized with digoxigenin-labeled probes against mouse Dlx1, Islet1, and Dlx1 targeting exon 4. The quantification of Dlx1/Satb2 double-labeled neurons was performed using ImageJ software (NIH, Bethesda, MD).
mice (n = 3 for each group). The average optical density (OD) of the NPY ISH signal was measured by Image Pro-Plus 6.0 (n = 3 for each group). Origin8 software was used for statistical analysis. All data were tested for normal distribution and homogeneity of variance. Comparisons were performed using the two-tailed Student’s t-tests. Statistical significance is displayed as *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001.

DATA AVAILABILITY

All data generated in this study are included either in this article or in the Supplementary Information files.

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AUTHOR CONTRIBUTIONS
Q.Z., L.Z., Y.H. and P.M. performed experiments and analyzed data; L.Z., B.M., N.-N.S. and Y.-Q.D. designed research studies; L.Z., N.-N.S. and Y.-Q.D. wrote the manuscript.

COMPETING INTERESTS
The authors declare no competing interests.

ETHICS STATEMENT
Our studies did not include human participants, human data, or human tissue. The animal studies were approved by the Laboratory Animal Ethics Committee of Fudan University (DSF-2020-041).

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Correspondence and requests for materials should be addressed to Ning-Ning Song.

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