Control of obesity and glucose intolerance via building neural stem cells in the hypothalamus

Juxue Li1,2,3, Yizhe Tang1,2,3, Sudarshana Purkayastha1,2,3, Jingqi Yan1,2,3, Dongsheng Cai1,2,3,*

ABSTRACT

Neural stem cells (NSCs) were recently revealed to exist in the hypothalamus of adult mice. Here, following our observation showing that a partial loss of hypothalamic NSCs caused weight gain and glucose intolerance, we studied if NSCs-based cell therapy could be developed to control these disorders. While hypothalamus-implanted NSCs failed to survive in mice with obesity, NF-κB inhibition induced survival and neurogenesis of these cells, leading to effects in counteracting obesity and glucose intolerance. To generate an alternative cell source, we revealed that iPS-derived NSCs were converted into htNSCs by neuropeptide treatment. Of note, obesity condition potentiated the transfer of carotid artery-injected NSCs into the hypothalamus. These iPS-derived cells when engineered with NF-κB inhibition were also effective in reducing obesity and glucose intolerance, and neurogenesis towards POMCergic and GABAergic lineages was accountable. In conclusion, building NSCs in the hypothalamus represents a strategy for controlling obesity and glucose disorders.

Keywords Neural stem cells; iPS; Hypothalamus; NF-κB; Neuropeptide; Obesity; Glucose tolerance

1. INTRODUCTION

Feeding, body weight and glucose balance are regulated by the mediobasal hypothalamus (MBH) in the central nervous system (CNS), and this regulation is mediated importantly by melanocortin signals from the arcuate nucleus (ARC), in particular anorexigenic proopiomelanocortin (POMC) neurons and orexigenic agouti-related peptide (AGRP) neurons [1–6]. These two neuronal types are sensitively and dynamically affected by systemic hormones such as leptin and insulin which fluctuate according to feeding and fat mass conditions, and this process is physiologically critical for maintaining body weight homeostasis [1–7]. However, when chronically challenged under high-fat diet (HFD) condition, these neurons reduce the responsiveness to leptin and insulin, and these changes contribute to the mechanism of HFD-induced obesity and type-2 diabetes (T2D) [1–7]. In addition to these signaling changes, research during recent years has begun to show that hypothalamic neural organization and neuronal numbers are altered under chronic HFD feeding [8–10]. For example, it was revealed that prolonged HFD feeding leads to a fractional loss (~10%) in POMC neurons in the hypothalamus [10–12]. Intriguingly, POMC gene expression is known to be divergently regulated [13,14], and was shown to be present in neural precursors which give rise to various neuronal types [15]. In relation to this background, we have been interested in understanding if impaired hypothalamic neurogenesis might play a role in the development of HFD-induced obesity. In agreement with this idea, several evidences were recently documented to indicate that the hypothalamus of adult rodents has neurogenesis [16–21]. We showed that the MBH contain adult hypothalamic NSCs (htNSCs), and these cells appear to be involved in the hypothalamic control of metabolic physiology [11]. Herein, we extended to study if htNSCs could be developed to treat obesity and related glucose disorders, and if so, what technological approaches could be introduced to promote the practical employment of this technique.

2. RESULTS

2.1. Partial loss of htNSCs leads to weight gain and glucose intolerance

We recently showed that dietary obesity is associated with impaired survival and neurogenesis of htNSCs [11]. In this context, we studied if htNSCs could be developed to treat or prevent against obesity and related metabolic disorders. We first tested if a loss of endogenous htNSCs in mice could be sufficient to lead to these metabolic disorders. To do so, we generated a mouse model with adult-onset ablation of dividing Sox2-positive htNSCs in the MBH. Briefly, adult C57BL/6 mice received MBH injection of lentiviruses expressing Sox2 promoter-driven Herpes simplex virus type-1 thymidine kinase (Hsv1-TK), a kinase that works to convert a nontoxic nucleoside analog, ganciclovir (GCV), into the toxic product which acts as a chain terminator during DNA replication and therefore kills dividing Sox2-positive cells [22–25]. Injection of matched control lentiviruses was performed in control mice for comparison. All these mice were then chronically treated with GCV to...
induce Hsv1-TK/GCV-mediated ablation of Sox2-positive hNSCs, and as assessed in Figure S1A and B, 3-month GCV treatment resulted in a partial loss of hNSCs in the ARC. Using this model, we studied if such a loss of hNSCs could affect the population of POMC cells in the hypothalamus, since POMC neurons were recently shown to undergo constant although slow-speed turnover [10,11]. Data revealed that there was ~10% reduction in POMC cells over a 4-month follow-up period in our model (Figure S1C). Through physiological profiling, we observed that these mice displayed a collection of metabolic disorders, including glucose intolerance, hyperinsulinemia, increased food intake and moderate weight gain (Figure S2). Thus, these data indicate that adult hNSCs play a significant role in body weight and glucose control, supporting us to study if hNSCs therapy could be developed to control related diseases.

2.2. hNSCs fail to survive under HFD-altered hypothalamic microenvironment

Subsequently, we investigated if exogenous hNSCs could be implanted into the MBH of mice to control HFD-induced obesity and related glucose disorders. To do so, we employed a line of hNSCs which we recently established [11]; in order to track hNSCs-induced neurogenesis, GFP was introduced into this cell line via lentiviral infection (Figure 1A and B), similarly as we performed previously [11], and we also verified the stem cell identity of these cells using immunostaining for a NSC marker (Figure 1B). For hypothalamic implantation, we targeted the MBH for cell injection shown in Figure 1C, given that the MBH is the hypothalamic region that is enriched with hNSCs. However, when these cells were injected in mice which underwent the development of HFD-induced obesity, this trial was unsuccessful, because implanted hNSCs survived poorly in the hypothalamic microenvironment of these HFD-fed mice. As shown in Figure 1D and E, the survival rate of implanted hNSCs decreased at Day 7, and then became undetectable at Days 15–30 post-injection. In contrast to HFD-fed mice, much better survival rate of these cells was seen when we implanted them into the MBH of chow-fed normal mice (Figure 1E). This data further supported that the hypothalamic microenvironment is altered under the condition of chronic HFD feeding, which was hostile for the survival of hNSCs. Thus, it is necessary to engineer hNSCs to increase the hypothalamic survival of these cells, in order to evaluate the potential of hNSCs-based cell therapy in treating overnutrition-related diseases such as obesity and T2D.

2.3. NF-κB inhibition promotes survival and neurogenesis of implanted hNSCs

According to the literature, chronic HFD feeding induces hypothalamic inflammation [26,27], and the proinflammatory pathway consisting of nuclear factor-κB (NF-κB) and its upstream IκB kinase-β (IKKβ) is a key mediator that links hypothalamic inflammation to disease outcomes [28–32]. Because obesity-associated hypothalamic IKKβ/NF-κB activation was known to disrupt hNSCs [11], we asked if NF-κB inhibition could help in vivo survival of implanted hNSCs. To test this idea, we used a line of hNSCs that was engineered with NF-κB inhibition, generated via lentiviral induction of dominant-negative IκBα (DNIκBα-hNSCs) (Figure 1A and B), and hNSCs containing lentiviral GFP (Con-hNSCs) were used as matched controls. We noted that compared to control cells, DNIκBα-hNSCs showed more neural branches, while their stemness remained as verified with immunostaining (Figure 1B). We implanted these cells into the MBH of mice under HFD feeding condition. Strikingly, survival rate of implanted DNIκBα-hNSCs increased dramatically, as we observed that ~40% of these cells survived at Day 30, at which we did not detect Con-hNSCs. In addition to survival success, implanted DNIκBα-hNSCs migrated and generated projections, implicating that these cells underwent neurogenesis and interactions with host hypothalamic cells. Then, we directly examined if NF-κB inhibition could promote neural differentiation of implanted hNSCs. Data revealed that while some implanted cells maintained stem cell identity (Figure 2A), 16% of these cells became to express NeuN (Figure 2B) at ~1 month post-implantation. Immunostaining data further revealed that a number of DNIκBα-hNSCs differentiated into cells which expressed POMC in chow-fed mice, and this effect was preserved under HFD feeding (Figure 2C and D). Thus, while chronic HFD feeding led to a loss in hypothalamic POMC cells, just as consistently shown in recent literature [10–12], this loss was ameliorated by DNIκBα-hNSCs-induced neurogenesis (Figure 2D). In addition, we found that leptin signaling was induced in some of DNIκBα-hNSCs-derived neural cells (Figure S3A), resulting in a restoration of leptin-induced feeding control in HFD-fed mice (Figure S3B). Altogether, NF-κB inhibition represents a solution for hNSCs to survive from HFD feeding-altered hypothalamic microenvironment.

2.4. Implanting NF-κB-inhibited hNSCs counteracts obesity and glucose disorder

While physiological implications of hNSCs implantation could be multiple, here we studied if it could exert an effect in counteracting HFD-induced metabolic disorders. For comparison, we included normal chow-fed mice as a dietary control in the study, and found that hypothalamic implantation of DNIκBα-hNSCs or Con-hNSCs did not affect the normal metabolic profiles of these mice, including body weight (Figure 2E), food intake (Figure 2F) and blood insulin levels (Figure S6C). Importantly, implantation of DNIκBα-hNSCs exerted significant effects in reducing body weight (Figure 2E), food intake (Figure 2F), glucose intolerance (Figure 2G) and blood insulin levels (Figure S4C) in HFD-fed mice. Body weight reduction in mice by DNIκBα-hNSCs was attributed specifically to fat loss (Figure S4A and B), further supporting the conclusion that the treatment provided an anti-obesity effect. In contrast, these metabolic effects were not produced by implanting Con-hNSCs (Figures 2E–G and S4C), which is consistent with the fact that these cells did not survive in HFD-fed mice. We also included vehicle injection in HFD-fed mice to provide a basal control, and data showed that the metabolic effects of injecting vehicle and Con-hNSCs were both negligible (Figure S5), again showing that NF-κB inhibition is necessary for the observed effects of hNSCs-based therapy. Overall, these effects of DNIκBα-hNSCs aligned with the data showing that implanted DNIκBα-hNSCs amended the loss of POMC cells in the hypothalamus in HFD-fed mice (Figure 2D). And these effects were also in agreement with the observation demonstrating that some of DNIκBα-hNSCs-derived cells responded to leptin (Figure S3). Based on these data, we concluded that introduction of hNSCs can work as a strategy for controlling obesity and related glucose disorders, and NF-κB inhibition is critical for success of this strategy.

2.5. Development of hNSCs phenotype by using iPS-derived NSCs

One of important goals in this research was to develop a strategy of hNSCs-based cell therapy that can be applicable for human diseases, and since it is unrealistic to collect endogenous human hNSCs, we explored if we could employ inducible pluripotent stem cells (iPS), a model of re-programmed stem cells [33]. Using a protocol established in the literature [34], we successfully induced NSCs from a line of mouse iPS, referred to as NSCs iPS (Figure 3A), as verified by our data showing that NSCs iPS expressed NSC markers Sox2 and nestin (Figure 3B). Also, just like native hNSCs, NSCs iPS differentiated into three neural lineages, including TuJ1-expressing neurons, GFAP-expressing astrocytes
and O4-expressing oligodendrocytes (Figure 3C). Interestingly, we found that implanted NSCsiPS in the MBH not only underwent neurogenesis (Figure S6A) but led to generation of POMC cells (Figure S6B). We speculated that hypothalamic microenvironment might provide necessary molecular stimuli to program NSCsiPS into htNSCs. To test this idea, we studied if treatment with hypothalamic neuropeptides could induce the differentiation of NSCsiPS into htNSCs. As established in the literature [35], ventral hypothalamic neurons are derived from embryonic NSCs, a process that is directed by hypothalamic programming involving gene upregulation of *Nkx2.1*, *Six3* and *Vax1*, and downregulation of *Pax6* in NSCs. Indeed, we observed that native htNSCs were characterized by increased levels in *Nkx2.1*, *Six3* and *Vax1*, but decreased levels in *Pax6* and *Rax* (Figure 3D). Intriguingly, this pattern of gene expression in htNSCs was reproduced in NSCsiPS through treatment with neuropeptide Y (NPY), but not other hypothalamic neuropeptides like *α*-MSH or cocaine/amphetamine-regulated transcript (CART). We also examined brain-derived

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**Figure 1:** Survival of hypothalamic implanted htNSCs engineered with NF-κB inhibition. Cell lines of Con-htNSCs (Con) and DNkBa-htNSCs (IκBα) were generated through lentiviral induction (A), examined for stemness and morphology via GFP and immunostaining of a NSC marker (B), and injected bilaterally in the MBH of HFD-fed vs. chow-fed male C57BL/6 mice (C–E). Images show technical success of delivering cells in the MBH at Day 3 post-injection (C), and longitudinal follow-up of indicated htNSCs in the MBH of HFD-fed mice (D). DAPI staining reveals nuclei of all cells in the sections. Scale bar=50 μm. Curves show longitudinal survival of htNSCs in HFD- vs. chow-fed mice (E). *P* < 0.05, **P** < 0.01 (compared to matched time points in the group of Con, chow), n=4–6 mice per group. Error bars reflect means ± s.e.m.
neurotrphic factor (BDNF), a neuropeptide that can promote brain neurogenesis [36], and found that conversion of NSCsiPS into htNSCs was further promoted by NPY and BDNF co-treatment (Figure 3D). Using immunostaining, we confirmed that while NPY treatment increased differentiation of NSCs iPS into Six3-expressing and Vax1-expressing cells, these effects increased when BDNF was added to the treatment (Figure 3E). Finally, since a fundamental role of hypothalamic neurogenesis is to generate cells that produce hypothalamic neuropeptides, we examined if this potential could be induced in the model NPY/BDNF-treated NSCsiPS. Data showed that, compared to native htNSCs, which can indeed be differentiated to produce *Agrp, Pomc* and *Cart* mRNAs, these changes were similarly produced in NPY/BDNF-treated NSCs iPS.

Figure 2: Neurogenesis and metabolic effects of implanting NF-κB-inhibited htNSCs. Dissociated NF-κB-inhibited htNSCs (IkBα) vs. Con-hNSCs (Con) were injected in the MBH of chow- vs. HFD-fed C57BL/6 mice, examined for stemness and neurogenesis of htNSCs at ~1 month post-injection via immunostaining of Sox2 (A–C), NeuN (B) and α-MSH (C), and effects of injected htNSCs on metabolic physiology (E–G). Histological data in A–D show MBH sections from HFD-fed mice injected with IkBα-hNSCs (A–C), and counting of endogenous (white bars) vs. exogenous hNSCs-derived (black, blue, red and green bars) POMC cells (D). Bar=50 μm. Physiological data show body weight (E), food intake (F) and GTT (G) obtained at indicated days (E) or 2–3 months post-implantation (F, G). *P<0.05, **P<0.01, ***P<0.001, compared between indicated exogenous cells (D) or compared to HFD-fed controls at matched time points (E, G), n=4 (D) and 5–6 (E–G) per group. Error bars reflect means ± s.e.m.
Figure 3: NSCs derived from iPS and conversion into htNSCs phenotype. Model of iPS-derived NSCs (NSCs iPS) was generated from iPS-derived embryoid bodies (A), verified via neurosphere staining of Sox2 and nestin (B), and assessed for differentiation via Tuj-1, GFAP and O4 immunostaining (C) and programming under neuropeptide treatment (D–F). D–F: NSCs iPS treated with neuropeptides including α-MSH, CART, NPY, or NPY and BDNF (NPY+B) were examined for induction of htNSCs-specific markers via quantitative PCR (D) and immunostaining (E), or subjected to differentiation followed by measurement of hypothalamic neuropeptide gene expression (F). Vehicle treatment was used to provide baseline levels profiles of various parameters. Native htNSCs under basal condition were included to provide positive controls. E: Bar = 50 μm, and Six3- and Vax1-positive cells were counted in lower panels. *P < 0.05, **P < 0.01, ***P < 10^-7, n=4 per group. Error bars reflect means ± s.e.m.
with DNI NSCsiPS via carotid artery, and for the control, we injected Con-NSCsiPS.

2.6. Obesity potentiates MBH penetration of carotid artery-injected NSCs

Following the in vitro studies in Figure 3, we tested if iPS-derived NSCs could be used to induce hypothalamic neurogenesis; however, just like htNSCs, NSCsPS also showed poor survival when injected in the MBH of mice with HFD-induced obesity. We then examined if this defect could be prevented when NF-κB inhibition was induced in these cells, and to do so, we generated a line of NSCsPS with lentiviral induction of DnkBx (GFP-conjugated), referred to as DnkBx-NSCsPS, and the control cell line was generated through lentiviral induction of GFP, termed Con-NSCsPS. Both of these cell lines were confirmed for stemness through immunostaining (Figure 4A), and we also noted that DnkBx-NSCsPS displayed more neural branches than control cells. Using this model, we studied if DnkBx-NSCsPS could be delivered to treat obesity in HFD-fed mice, and to increase the application value, we studied if carotid artery injection could be used instead, since this method has been adopted in mice, and to increase the application value, we studied if carotid artery injection could be used instead, since this method has been adopted in

2.7. Therapeutic effects of carotid artery-delivered NSCs

As described in Figure 4, HFD-fed mice were injected with DnkBx-NSCsPS via carotid artery, and for the control, we injected Con-NSCsPS in matched HFD-fed mice. In the control group, Con-NSCsPS failed to survive, and thus as expected, we did not see an anti-obesity effect. However, delivery of DnkBx-NSCsPS led to ~30% decrease in HFD intake (Figure 5A), and this feeding control was associated with a moderate effect in weight reduction which lasted although only a short period (Figure 5B). When mice continued to be fed on the HFD for long period, the anti-obesity therapeutic effect of DnkBx-NSCsPS declined and diminished; however, the extent of weight gain in these mice under HFD feeding was much less substantial, compared to the levels of weight gain in control mice (Figure 5B). We also examined if hypothalamic leptin resistance, a key event in HFD-induced obesity and T2D, could be reduced by this cell therapy in mice. Data showed that while control mice showed impaired action of leptin in suppressing food intake over 24-h experimental period, the effect of leptin in controlling feeding was significantly restored in mice treated with DnkBx-NSCsPS (Figure 5C). Furthermore, consistent with the effects on feeding and weight control, treatment with DnkBx-NSCsPS led to improved glucose tolerance in HFD-fed mice (Figure 5D). In summary, NSCsPS can be developed as a practical source of cells for controlling obesity and glucose disorders, and artery injection as a delivery approach.

2.8. Metabolic effects of NSCs mechanistically via neurogenesis

Supported by the potential application value of NSCsPS, we further studied how these cells could work to generate metabolic effects in mice. In addition to the actions of these cells in inducing neuropeptides (Figure 3F), we investigated if neurotransmitter pathway(s) could be introduced by neurogenesis of these cells. Thus, for the experiment described in Figure 3F, we further analyzed molecules in GABAAergic and glutamatergic pathways, including glutamic acid decarboxylase 67 (GAD67) and GAD65 (encoded by Gad1 and Gad2, respectively), vesicular GABA transporter (vGAT, encoded by Slc32a1), vesicular glutamate transporters 2 (vGLUT2, encoded by Slc17a6), and gial glutamate transporter 1 (GLT-1, encoded by Slc1a2). First, using differentiated hNSCs as positive control, we found a pattern of changes in GABAAergic pathway, including increases in Gad1 and Slc32a1 mRNA levels but a decrease in Gad2 mRNA levels (Figure 6A). Molecules in glutamatergic pathway also changed, including Slc1a2 and Slc17a6 mRNA levels, but to a lesser degree compared to changes in Slc32a1 mRNA levels (Figure 6A). Interestingly, when treated with NPY and BDNF, differentiated NSCsPS showed similar molecular changes in GABAAergic and glutamatergic pathways (Figure 6A). In this background, we then asked if the metabolic effects of DnkBx-NSCsPS were attributed to a neurepoxide or a neurotransmitter or both. To address this question, we designed a “rescue” experiment that is illustrated in Figure 6B; in this study, we used lentiviral RNAi to knockdown various neuropeptides or neurotransmitter-regulatory molecules in DnkBx-NSCsPS, and then implanted these cells into the MBH of HFD-fed mice. Data revealed that the metabolic effects of DnkBx-NSCsPS in feeding, body weight and glucose tolerance were abrogated by ablation of either Pomc or Gad67 (Figure 6C–E). Therefore, NSCs-derived neural cells can lead to at least two different types of differentiated neural cells which express Pomc and Gad67, respectively, and these cell types can work together to co-mediate the therapeutic effects of NSCs (Figure 6F), a model which is in agreement with the understanding [40] that GABAAergic neurons and POMC neurons may have independent actions in regulating body weight.

3. DISCUSSION

Adult NSCs are a small population of cells with slow dividing rate in the brain [41], and because of this feature, it remains largely unclear regarding the potential physiological role of these cells. More recent in vitro and in vivo studies suggested that the hypothalamus is a brain region that contains adult NSCs [10,11,17,18,42]. In the current work, using site-specific ablation of hNSCs in the mediobasal hypothalamic region of adult mice, we found that these animals developed overeating, weight gain and glucose intolerance, a collection of disorders which can be similarly induced by chronic HFD feeding. As this defect of hNSCs was recently revealed as a neurodegenerative basis for obesity and pre-diabetes [10–12], hNSCs-based cell therapy may have a potential as a long-term solution for these metabolic diseases, which can fix the involved central dysregulation in a comprehensive manner which may not be offered by a drug chemical which often targets a specific signaling cascade. Indeed, recent research in neuroscience has begun to suggest that NSCs have therapeutic potential for treating neurological diseases [43–47]. In this background, here we studied if hNSCs-based therapy could be developed to control obesity and metabolic diseases.
However, we found that hypothalamic microenvironment of obesity condition is hostile for implanted htNSCs, and this condition leads to poor survival and neurogenesis of these cells. To solve this problem, we discovered that engineering of NSCs with NF-\(\kappa\)B inhibition renders implanted htNSCs to survive and undergo neurogenesis under obesity condition. This success is related to the fact that obesity is closely associated with hypothalamic inflammation [26,27], and NF-\(\kappa\)B inhibition is strongly counter-inflammatory and thereby protects implanted htNSCs from being disrupted by hypothalamic inflammation [11]. According to our recent understandings, we also reasoned that NF-\(\kappa\)B in neurogenesis engages a programmatic switch from neuronal genesis to gliogenesis, a process which can be similarly induced by Notch pathway [48], and agreeably, our recent work has demonstrated that Notch pathway is responsible for the role of NF-\(\kappa\)B in NSCs-induced neurogenesis [11]. Interestingly, obesity-related hypothalamic inflammation was recently shown to be associated with gliosis [12], and thus, if NF-\(\kappa\)B-inhibited htNSCs can be provided to mediate programmatic switch, this treatment may lead to suppression in gliosis and therefore improvement of hypothalamic functions from another biological perspective.

In this research, we also considered that it is unrealistic to obtain human htNSCs for an application. To address this limitation, we explored if iPSC could be used as an alternative source for generating NSCs. Indeed, we successfully derived NSCs from iPSC, and further found that conversion of iPSC-derived NSCs into htNSCs was induced by treatment with NPY and BDNF. Moreover, survival and neurogenesis of iPSC-derived NSCs were improved by NF-\(\kappa\)B inhibition to induce protective effects against obesity and glucose intolerance. Hence, iPSC-derived NSCs have a potential for

Figure 4: Artery delivery of NF-\(\kappa\)B-inhibited NSC\(^{\text{ps}}\) and in vivo neurogenesis. Lines of DN\(\kappa\)B\(\alpha\)-NSC\(^{\text{ps}}\) and Con-NSC\(^{\text{ps}}\) were verified via GFP and immunostaining of a NSC marker (A), and delivered into HFD-fed mice via carotid artery injection (B–E). Brain sections were prepared to analyze the hypothalamic transfer of artery-injected DN\(\kappa\)B\(\alpha\)-NSC\(^{\text{ps}}\) at ~1 month post-injection (C) and hypothalamic neurogenesis at ~3 months post-injection via immunostaining of NeuN (C) and POMC (D, E). Bar graphs show counting of endogenous cells (white bars) vs. exogenous NSC\(^{\text{ps}}\)-derived cells (black, blue, red and green bars) that expressed POMC (E). \(^*P<0.05\), \(^{**}P<0.01\), \(^{***}P<0.001\), (comparisons between the columns of exogenous cells), n=4 mice per group. Error bars reflect means ± s.e.m. Scale bar=50 \(\mu\)m (A, B) and 25 \(\mu\)m (C).
being developed as a source of hNSCs to treat or control metabolic diseases. In addition to solving cell source issue, we further considered that the method of hypothalamic implantation is not practical for human application, and knowing that the BBB in the MBH becomes penetrable under obesity/diabetic conditions [12,39], we performed an initial study to test if NSCs could be delivered via carotid artery to exert a therapeutic effect, and the overall findings were supportive. Nevertheless, we recognized that the anti-obesity therapeutic effect which we obtained was rather short-term, indicating that it is necessary to optimize the protocol, perhaps even in combination with other approaches, in order to increase the therapeutic efficacy. In light of the potential mechanism for the metabolic effects of NSCs therapy, we provided data showing that neurogenesis of these cells towards POMCergic and GABAergic lineages was at least accountable. On the

Figure 5: Metabolic effects from artery injection of NF-κB-inhibited NSC iPS. Male C57BL/6 mice were induced to develop dietary obesity through 3-month HFD feeding, and then received intra-carotid artery injection of \(2 \times 10^5\) DNI\(\alpha\)-NSCiPS (I\(\kappa\B alpha\)iPS, Con) or Con-NSC iPS (Con iPS, Con). Mice continued to be maintained on the HFD for the entire experiment. Data show food intake (A), body weight (B), and leptin-induced feeding inhibition in response to an i.c.v. injection of 5 μg leptin (Lep, L) vs. vehicle (Veh, V) following 6-h fasting (C), and blood glucose levels during GTT (D). Data were obtained at the indicated weeks (W) (A, B and D), or 15 weeks (C) post-surgical recovery. * \(P<0.05\), ** \(P<0.01\), \(n=5\) mice per group (A, B and D), and \(n=4\) mice per group (C). Error bars reflect means ± s.e.m.
other hand, we also want to point out that many of these NSCs-derived cells may represent neuronal precursors rather than mature neurons, and it is possible that the observed metabolic effects of NSCs therapy were mostly attributed to the secretory factors rather than synaptic actions of their derived neural cells. In conclusion, despite many limitations which remain to be addressed, data which we presented can lend a rationale and biotechnological basis for future interest in developing NSCs to control obesity and related diseases.

4. METHODS

4.1. Lentiviruses and hTNSCs

The cDNAs for \( \text{IkB}^\alpha \) and control GFP have been described previously \([11,26,30,49]\). Lentiviral vectors containing POMC shRNA, GAD67 shRNA or scramble shRNA were obtained from Sigma. Lentiviruses were produced by co-transfecting viral vectors with the package plasmids into HEK 293 FT cells, as previously described \([11,26,49]\). Lines of hTNSCs

![Figure 6](image-url)
were generated according to our protocol described previously [11]. Briefly, the hypothalamus was dissected from mice and cut into small pieces (~1 mm³), and digested with 0.25% papain (Worthington) for 30 min at 30 °C and gently triturated approximately 10 times using fire-polished tips. The desired cell population was suspended in growth medium containing Neurobasal-A (Invitrogen), 2% B27 without vitamin A (Invitrogen), 10 ng/ml EGF (Sigma) and 10 ng/ml bFGF (Invitrogen), seeded in ultralow adhesion 6-well plates (Corning) at a density of 100,000 cells per well and incubated in 5% CO₂ at 37 °C. On day 7, neurospheres were collected through low-speed centrifugation, dissociated into single cells by trypsinization using TrypLE express media, and processed for passaging or cryopreservation. Lentiviruses-induced lines of hNSCs were prepared by subjecting hNSCs in growth medium with lentiviruses for 3 days, followed by selection process through addition nucleoside antibiotic blasticidin (1 μg/ml) to culture medium. Cells transduced with lentiviruses were maintained in blasticidin-containing selection growth medium over several passages until gene knockdown was stably established.

4.2. Pluripotent stem cells (iPS) and derived NSCs

Stemgent™ Mouse Primary iPS Cells-NNeo was purchased from STEMGENE Company. iPS were maintained in standard embryonic stem cell culture conditions. Briefly, irradiated mouse embryonic fibroblast were plated at a density of 2.5 × 10⁴ cells per cm² as feeder cells in gelatin-coated 6-well plates, and iPS were maintained on the feeder cells with standard embryonic stem cells culture medium containing knock-out DMEM, 10% knock-out serum, 2 mM L-Glutamine, 0.1 mM nonessential amino acids, 1 mM sodium pyruvate, 0.1 mM β-mercaptoethanol and 10 ng/ml LIF (all from Invitrogen). For embryoid body (EB) formation, dissociated iPS cells (via 0.05% trypsin-EDTA solution) were cultured at 5 × 10⁴ cells per ml in the EB formation medium (standard embryonic stem cells culture medium without LIF). Following 3 days of EB formation, cultured cells were stimulated with 5 μM RA (Sigma) for 7 days with culture medium changed every day. At Day 11, EBs were dissociated into single cells and transfer to NSC culture medium (the growth medium described above). Spheres formed after 2 passages of culture were mainly neurospheres, as confirmed by immunostaining of NSC markers, and were also examined for multi-potent differentiation into 3 neural cell lineages. To generate iPS-derived NSC with stable gene manipulation, dissociated neurospheric cells were infected with various lentiviruses (containing fluorescent marker GFP), and selected over passages using the blasticidin-containing selection medium. Neuropeptide treatment: iPS-derived NSCs were cultured in NSC culture medium in the presence of α-MSH (20 ng/ml), CART (20 ng/ml), NPY (20 ng/ml) or BDNF (20 ng/ml) either individually or in combination for 7 days, with medium replaced every 2 to 3 days. Cells were collected for real-time PCR analysis or immunofluorescence microscopy.

4.3. MBH implantation

Bilateral injections of mediobasal hypothalamus were performed in adult C56BL/6 mice as described previously [26,30,49]. Briefly, under an ultra-precise stereotax, cells at the number of 8000 suspended in 0.5 μl PBS was injected into each side of the mediobasal hypothalamus via guide cannula directed to the coordinates of 0.17 cm posterior to the bregma, 0.03 cm lateral to the middle line, and 0.50 cm below the skull surface of mice.

4.4. Carotid artery injection

C57BL/6 mice were anesthetized and a longitudinal incision is made on the right side of the neck. Under a dissecting microscope, the common, internal, and external carotid arteries are dissected, and the external carotid artery and the proximal part of the common carotid artery were clamped by a microclamp. Cell injections were performed by a 33-gauge microneedle punctured into common carotid artery. For each injection, 2 × 10⁵ cells suspended in 100-μl PBS are injected with a preserved flow. After cell injection, needle was withdrawn, and a pressure in the injection site was maintained until bleeding stopped. After surgical procedure, mice were kept in a warm environment until recovery.

4.5. Metabolic phenotyping

C57BL/6 mice were obtained from Jackson Laboratory, and were housed in standard conditions. High-fat diet was obtained from Research Diets, Inc. Body weight of individually housed mice was measured twice per week and food intake was recorded on daily basis. MRI measurement of lean vs. fat mass was performed at the core facility at Albert Einstein College of Medicine. For GTT, overnight fasted mice were injected with glucose (2 g kg⁻¹ body weight) intraperitoneally, and blood glucose levels at various time points were measured using a Glucometer (Bayer). Fasting blood insulin levels were measured using serum or plasma samples obtained from mice following 14 to 20 h of fasting. All procedures were approved by the Institutional Animal Care and Use Committee of Albert Einstein College of Medicine.

4.6. Immunostaining

For tissue sections, mice under anesthesia were trans-heart perfused with 4% PFA, and the brains were removed, post-fixed in 4% PFA for 4 h, and infiltrated in 20–30% sucrose. Brain sections (20 μm) were made using a cryostat at −20 °C. Cultured cells on coverslips were fixed with 4% PFA for 10 min at room temperature. For immunostaining of culture cells, cells were prepared onto cover slides and fixed with 4% PFA. Fixed tissue sections/cells were blocked with serum of appropriate species, penetrated with 0.2% Triton-X 100, treated with primary antibodies (Millipore), mouse anti-Sox2 antibody (R&D Systems), sheep anti-POMC (Phoenix), mouse anti-GFAP, anti-O4, anti-NeuN (Millipore), rabbit anti-AGRP (Phoenix Pharmaceuticals), mouse anti-GFAP, anti-Stat3 (Cell Signaling), Rabbit anti-Six3 and anti-Vax1 antibodies were kindly provided by Guillermo Oliver and Greg Lemke, respectively. DAPI (Vector) staining revealed the nuclei of all cells in the slides. Images of immunostaining were captured under a confocal microscope. Cell counting of immunostaining: serial hypothalamus sections across the MBH were made using a cryostat at −20°C. Cultured cells on coverslips were fixed with 4% PFA for 10 min at room temperature. For immunostaining of culture cells, cells were prepared onto cover slides and fixed with 4% PFA. Fixed tissue sections/cells were blocked with serum of appropriate species, penetrated with 0.2% Triton-X 100, treated with primary antibodies and followed by reaction with Alexa Fluor™ 488 or 555 or 633 secondary antibodies (Invitrogen). Naïve IgGs of appropriate species were used as negative controls. Primary antibodies included rabbit anti-POMC (Phoenix Pharmaceuticals), mouse anti-GLP, anti-04, anti-NeuN antibodies (Millipore), mouse anti-Sox2 antibody (R&D Systems), sheep anti-α-MSH (Millipore), rabbit anti-POMC (Phoenix), rabbit anti-AGRP (Phoenix), and rabbit anti-pSTAT3 (Cell Signaling). Rabbit anti-Six3 and anti-Vax1 antibodies were kindly provided by Guillermo Oliver and Greg Lemke, respectively. DAPI (Vector) staining revealed the nuclei of all cells in the slides. Images of immunostaining were captured under a confocal microscope. Cell counting of immunostaining: serial hypothalamus sections across the MBH were made at the thickness of single cell (10 μm), and every 5 sections were represented by one section with staining and cell counting. The numbers in representative sections were multiplied by 5 to indicate the total numbers. Cell counting in cultured cell slides was performed by including at least 4 to 6 random fields per slide under microscopy.

4.7. Quantitative RT-PCR

Total RNA was extracted using TRIzol (Invitrogen) following the manual. cDNA was synthesized using the M-MLV reverse transcriptase (Promega). Real-time PCR was performed using the StepOnePlus™ Real-Time PCR System and SYBR Green PCR Master Mix (Applied Biosystems). Relative gene expression levels were normalized against the mRNA level of the house-keeping gene β-actin.
4.8. Statistical analyses

Kolmogorov–Smirnov test was used to determine parametric distribution of data sets. ANOVA and appropriate post-hoc analyses were used for comparisons involving more than two groups, and two-tailed Student’s t tests were used for comparisons involving only two groups. Data are presented as mean ± s.e.m. P < 0.05 was considered significant.

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CONFLICT OF INTEREST

The authors state that they have no competing financial interests.

APPENDIX A. SUPPLEMENTARY MATERIALS

Supplementary data associated with this article can be found in the online version at http://dx.doi.org/10.1016/j.molmet.2014.01.012.

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