Endoplasmic reticulum stress contributes to the decline in doublecortin expression in the immature neurons of mice with long-term obesity

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Adult hippocampal neurogenesis (AHN) plays an important role in hippocampus-dependent function. The number of doublecortin (Dcx)-positive immature neurons in the dentate gyrus decreases over time, especially in the early stages of Alzheimer’s disease (AD), and is further reduced in later stages of AD. Obesity in midlife is associated with dementia later in life; however, the underlying mechanisms by which obesity results in the development of dementia later in life remain unknown. Here, we show that endoplasmic reticulum (ER) stress was activated in the hippocampus and processes of Dcx-expressing immature neurons were shortened, coexpressing CHOP in APP23 AD model mice with high-fat diet-induced long-term obesity and in aged Leprdb/db (db/db) mice. Moreover, in cells differentiating from hippocampal neurospheres, Dcx mRNA was rapidly degraded via a microRNA (miRNA) pathway after thapsigargin treatment in vitro. These results indicate that loss of Dcx mRNA induced by ER stress during AHN may cause memory impairment in obese individuals later in life.

Abbreviations
ATF4  Activating transcription factor 4
Aβ  Amyloid-β
eIF2α  Eukaryotic translation initiation factor 2α
ER  Endoplasmic reticulum
PBS  Phosphate-buffered saline
PCR  Polymerase chain reaction
siRNA  Small interfering RNA
UTR  Untranslated region
PKR  Protein kinase RNA-activated
PERK  PKR-like ER-localized eIF2α kinase
GCN2  General control nonderepressible 2
IRE1  Inositol-requiring protein-1
CHOP  C/EBP-homologous protein

New neurons are generated in distinct regions of the adult human brain, i.e., the hippocampal subgranular zone (SGZ) and subventricular zone (SVZ)¹. A total of 700 new neurons are generated daily in the hippocampus, but this number declines during aging². Immature neurons in the dentate gyrus (DG) were not detected beyond adolescence in available human hippocampi³,⁴. However, under physiological conditions, adult hippocampal neurogenesis (AHN) was recently shown to be involved in hippocampus-dependent functions such as pattern separation⁵ and stress resilience⁶, and AHN is impaired in patients with Alzheimer’s disease (AD)⁷ and depression⁸. The number of cells positive for doublecortin (Dcx), which is specifically expressed in immature neurons, in the DG is decreased in patients with mild cognitive impairment (MCI)⁹ and those in the early stages of AD. Obesity in midlife is associated with dementia later in life; however, the underlying mechanisms by which obesity results in the development of dementia later in life remain unknown. Here, we show that endoplasmic reticulum (ER) stress was activated in the hippocampus and processes of Dcx-expressing immature neurons were shortened, coexpressing CHOP in APP23 AD model mice with high-fat diet-induced long-term obesity and in aged Leprdb/db (db/db) mice. Moreover, in cells differentiating from hippocampal neurospheres, Dcx mRNA was rapidly degraded via a microRNA (miRNA) pathway after thapsigargin treatment in vitro. These results indicate that loss of Dcx mRNA induced by ER stress during AHN may cause memory impairment in obese individuals later in life.
of AD and is further reduced in later stages. Staining for markers of various differentiation stages has revealed substantial impairment of Dcx-positive cell maturation in the DG of subjects with AD. Dcx, which stabilizes and maintains microtubules in maturing neurons, has been identified as a causal gene of X-linked lissencephaly and double cortex syndrome, resulting in the arrest of neuronal migration and a long 3′ untranslated region (UTR), indicating its involvement in the regulation of mRNA stability and suggesting that Dcx mRNA is a target of microRNA (miRNA)-128. Consistently, the Dcx transcript level is increased in Dicer−/− neural stem cells (NSCs) female mice heterozygous for a mutation in Dcx show abnormalities in hippocampal lamination, defects in contextual and conditioned fear memory, and mild deficits in Morris water maze (MWM) performance, indicating that Dcx expression in the hippocampus is important for learning.

The number of people living with dementia is increasing; however, age-specific incidence rates of dementia have recently decreased in several countries because of decreases in the prevalence of some risk factors due to lifestyle changes. Twelve modifiable risk factors are known, including obesity, and eliminating these risk factors could prevent or delay the onset of dementia. Consistently, the memory abilities of AD model mice with high-fat diet (HFD)-induced obesity and of the offspring of AD model mice crossed with diabetic ob/ob mice or diabetic db/db mice are impaired compared with that of AD model mice. A 28-year study on changes in body mass index (BMI) revealed that obesity at age 50 is associated with dementia, indicating that obesity in midlife is related to dementia later in life. However, the underlying mechanisms by which obesity results in the development of dementia later in life remain unknown. Interestingly, the survival and proliferation of hypothalamic NSCs are disrupted by IκB kinase β (IKKβ)/nuclear factor-xB (NF-xB) activation in the context of long-term HFD-induced obesity, and a decline in the release of exosomal miRNA due to the loss of hypothalamic NSCs regulates aging. Several studies have suggested that endoplasmic reticulum (ER) stress is critical for obesity-induced inflammation. Analysis of the brain transcriptomes of mice with different APOE genotypes and previously published RNA sequencing (RNA-seq) data from the hippocampus and superior frontal cortex subjects has shown that the expression of its involvement in transcription factor 4 (ATF4), which is increased during ER stress during AHN may underlie the memory impairment in obese individuals later in life.

Results
Behavior to novel object and swimming capacity were compromised in AD model mice with long-term obesity and diabetes model mice. We previously showed that the expression levels of ATF4 are increased in the cortex, hippocampus, and amygdala in the offspring of obese and diabetic APP23 AD model mice crossed with db/db mice through a process known as the ISR. ATF4 expression in the brain is decreased by quercetin, and this change is accompanied by an improvement in memory. To examine the effect of ER stress on AHN in the brains of mice with long-term obesity, we first examined whether memory was impaired in HFD-induced obese mice. Obesity was induced by HFD feeding for 41 weeks in 66-week-old APP23 AD model mice (Fig. 1a–d) and for 64 weeks in 74-week-old wild-type C57BL6 mice (Suppl. Fig. 2). The serum levels of fasting blood sugar (FBS) and insulin were increased in both groups of mice subjected to long-term HFD feeding compared with mice fed a standard diet (Fig. 1b and Suppl. Fig. 2b). To examine whether long-term HFD-induced obesity affects memory, we performed the novel object location (NOL) test. Analysis of the ratio of the time spent exploring the target object to the time spent exploring all three objects revealed that the cognitive capacity of APP23 mice with long-term obesity was not significantly different from that of APP23 mice fed a standard diet. However, the time spent exploring the object at the novel location was significantly decreased in APP23 mice with long-term obesity, as they potentially exhibited neophobic behavior (Fig. 1c and d). Next, we examined whether spatial memory was affected in sixteen-week-old Lep db/db (db/db) mice, which had a higher body weight than wild-type and heterozygous littermates (control), even at 6 weeks of age (Suppl. Fig. 1a). In the MWM test, the latency to find the invisible platform during the acquisition phase was not changed and the number of platform crossings made by db/db mice was significantly decreased; these parameters were considerably affected by the differences in swimming speed and distance (Suppl. Fig. 1b). We next assessed aged wild-
type mice with long-term obesity (Suppl. Fig. 2). The index for the object at the novel location was significantly increased in mice fed a standard diet but not in mice with long-term obesity (Suppl. Fig. 2e). These results suggested that long-term obesity affect spatial memory and behavior in mice.

**ER stress was activated in the brains of mice with long-term obesity.** To investigate the role of ER stress in the brains of mice with long-term obesity, we first used 59- to 61-week-old APP23 mice. After eight weeks of HFD feeding beginning at 21 weeks of age, the body weights of the mice fed a HFD were significantly increased. The body weights of the mice fed a HFD and the mice fed the standard diet were 42.5 ± 0.9 g and 28.3 ± 3.6 g, respectively, when the mice were approximately 60 weeks of age. Western blot analysis showed that the expression levels of CHOP in the hippocampi of APP23 mice with long-term obesity were increased compared with those in the hippocampi of mice fed the standard diet (Fig. 2a). After four weeks of HFD feeding, obesity was detected in wild-type C57BL/6 mice at the age of 16 weeks, and the body weights of mice fed a
HFD and the standard diet were 55.8 ± 9.7 g and 36.0 ± 3.9 g, respectively, at the age of 86 weeks. Western blot analysis showed that the expression levels of CHOP in the hippocampi of C57BL6 mice with long-term obesity (a) and db/db mice (b), st.: standard diet; H: high-fat diet. The arrow in each panel indicates ATF4 or CHOP and the arrowhead in each panel indicates Dcx. The asterisk indicates nonspecific signal (b). (c) ATF6 and Xbp-1 expression in the hippocampi of 60-week-old db/db mice. The arrowheads in each panel indicate the ATF6 fragment or Xbp-1 s. ES: mouse embryonic stem cell line. ES cells were treated with 2 mM DTT for 1 min or 30 min. (d) Semiquantitative RT–PCR of Chop in the hippocampi of aged control and db/db mice. (a–c) Statistical significance (p < 0.05) was determined by Student’s t test. The error bars represent the SEMs. n.s.: not significant.
expression of Xbp-1s and ATF6 fragmentation in the hippocampi of 60-week-old db/db mice. We found that the N-terminal cleavage product of ATF6 and Xbp-1s were not detected in the hippocampi of aged db/db mice but were detected in mouse ES cells treated with 2 mM dithiothreitol (DTT) (Fig. 2c). Therefore, we examined the levels of Chop mRNA in the hippocampi of aged control and db/db mice. We found that Chop mRNA levels were not increased in db/db mice (Fig. 2d), suggesting that ATF6 is not involved in CHOP induction in aged db/db mice. Xbp-1s expression was not observed in aged db/db mice. These results suggested that long-term obesity induced ER stress, particularly the ISR, in the hippocampus.

CHOP and Dcx were coexpressed in the immature neurons of mice with long-term obesity. To investigate the impact of the ER stress on AHN, we performed immunostaining for Ki67, which is continuously expressed by proliferating neurons, and Dcx, a marker for immature neurons. The number of Ki67-positive cells in the hippocampal DGs of db/db mice was similar to that in the DGs of control mice in both the young and aged groups (Fig. 3a, b). Additionally, the levels of mature BDNF in the hippocampi of 45-week-old control and db/db mice were similarly measured (Fig. 3c). The levels of Dcx in APP23 mice fed a HFD were significantly decreased compared with those in APP23 mice fed a standard diet (Fig. 2a), but the levels of Dcx did not differ between C57BL6 mice fed a HFD and C57BL6 mice fed a standard diet or between/db/db and control mice (Fig. 2b and Suppl. Fig. 3). Therefore, we performed immunohistochemistry to examine the expression of Dcx in the hippocampus. Since Dcx is sensitive to postmortem breakdown, the mice were anesthetized and quickly perfused with 4% paraformaldehyde for fixation, and then the expression of Dcx in the hippocampi of control and db/db mice was then examined by immunohistochemistry. The number of Dcx-positive cells in the DG did not differ between 45-week-old control and db/db mice (Fig. 3d, f) or between 10-week-old control and db/db mice (data not shown). We observed reduction of Dcx expression in processes in z-stacks by confocal microscopy of Dcx-positive cells in the DG in 45-week-old db/db mice (Fig. 3e). Using the method described by Plump et al. for the characterization of Dcx-positive cells, more cells with short processes were observed in 45-week-old db/db mice than in 45-week-old control mice (Supplementary Fig. 4). These Dcx-positive cells in 45-week-old db/db mice expressed CHOP (Fig. 3g). Consistently, we observed that the Dcx-positive cells of APP23 mice with long-term obesity expressed CHOP (Fig. 3b). These results suggested that the ER stress was activated in the Dcx-positive immature neurons of mice with long-term obesity.

Thapsigargin reduced Dcx expression in immature neurons through Dcx mRNA degradation. To investigate the effect of the ER stress on the Dcx expression in immature neurons, we cultured neurospheres isolated from the mouse hippocampus according to a previously established protocol. After neurospheres were cultured on laminin-coated dishes for 5 days in vitro (DIV) (Fig. 4a), several marker proteins, i.e., Dcx, calreticulin, Nestin, and βIII tubulin, were expressed (Fig. 4c). Therefore, we challenged these cells with thapsigargin, a sarcoplasmic reticulum/ER Ca²⁺-ATPase inhibitor, for 6 h and then cultured them in new medium to induce ER stress according to a previously published method. The expression levels of Dcx corrected for GAPDH after thapsigargin treatment were much lower than those after DMSO treatment as determined by western blot (Fig. 4c); however, the differences were not significant as determined by Student’s t-test (p = 0.17). Consistently, immunostaining revealed that the processes in cells expressing both Dcx and CHOP were not decreased by thapsigargin treatment (Fig. 4d), as determined by western blotting, we measured the levels of Dcx mRNA by semiquantitative reverse transcriptase polymerase chain reaction (RT–PCR) (Fig. 4e). We found that the mRNA levels of Dcx, but not Nestin or p35, quickly decreased during incubation in new medium after thapsigargin treatment (lower panel in Fig. 4e). The levels of Dcx mRNA were not rescued by the IRE1 inhibitor 4µ8C, indicating that Dcx mRNA is not a substrate for RIDD (Fig. 4f). Since the Dcx transcript has a long 3’UTR, which binds the Musashi RNA-binding protein, mRNA stability and translation are possibly regulated by the binding proteins. We found that the levels of Dcx mRNA were not rescued by zVAD (Fig. 4g), suggesting that caspase substrates are not involved in Dcx mRNA stability and that apoptosis does not decrease Dcx mRNA expression. Next, we knocked down Dicer in differentiating NSCs using a small hairpin RNA (shRNA) and small interfering RNA (siRNA) and then treated the cells with thapsigargin to examine whether the ER stress-induced reduction in Dcx mRNA expression was regulated by Dicer, as Dcx mRNA expression has been reported to be upregulated in Dicer-deficient NSCs. Neurospheres cultured on laminin were infected with lentiviral particles carrying an shRNA targeting Dicer at 1 DIV, transfected with an siRNA targeting Dicer at 4 DIV, and then treated with thapsigargin at 5 DIV. Real-time RT–PCR showed that the thapsigargin-induced downregulation of Dcx mRNA expression was significantly prevented by the knockdown of Dicer in differentiating NSCs (Fig. 4h).

To identify the miRNAs upregulated during ER stress, we performed small RNA-seq on differentiating neurospheres. Analysis of the differential expression of microRNAs in thapsigargin-treated immature neurons revealed that some microRNA, such as miR-148a-5p, miR-129b-3p, and miR-135a-2-3p, were significantly increased upon thapsigargin treatment (Fig. 5a).
Figure 3. CHOP and Dcx were coexpressed in the immature neurons of mice with long-term obesity. (a, b) Ki67-positive (+) cells in the SGZ of the DG/section in 10-week-old (control mice, n = 2; db/db mice, n = 2) (a) and 45-week-old (control mice, n = 2; db/db mice, n = 2) (b) db/db mice. Scale bars: 100 µm. The box-and-whisker plot represents the number of cells per section from each animal. Statistical significance (p < 0.05) was determined by one-way ANOVA followed by the Bonferroni post-hoc test. (c) Mature BDNF expression in the hippocampus of 46-week-old mice. Control mice (n = 5); db/db mice (n = 5). Statistical significance (p < 0.05) was determined by Student’s t test. (d-f) Number of Dcx+ cells in the DG/section in 45-week-old control and db/db mice. Serial sections of the DG from zero to 6.30 µm imaged by confocal microscopy. Scale bars: 40 µm (d); 20 µm (e). The box-and-whisker plot represents the number of cell per section from each animal. Control mice (n = 2); db/db mice (n = 2) (f). Statistical significance (p < 0.05) was determined by one-way ANOVA followed by the Bonferroni post-hoc test. (g, h) CHOP was expressed in the Dcx-expressing neurons in 45-week-old db/db mice (g) and 69- to 71-week-old APP23 mice with long-term HFD-induced obesity (h). (g) Control mice (n = 2, 12 brain slices); db/db mice (n = 2, 12 brain slices). (h) Standard diet APP23 mice (n = 2, 8 brain slices); high fat diet APP23 mice (n = 3, 12 brain slices). The arrowheads in each panel indicate Dcx- and CHOP-positive cells. Scale bars: 20 µm. The fluorescence intensity profiles are shown in the square boxes (bottom panels).
Figure 4. Thapsigargin reduced Dcx expression in immature neurons by decreasing the Dcx mRNA levels. (a) Experimental schedule and representative images. Hippocampal neurospheres from mice were differentiated on laminin for 5 days. The cells were treated with thapsigargin (Tg) for 6 h, washed once with medium, and then cultured in medium without Tg for the indicated amounts of times. (b) Immunostaining for Dcx, CHOP, and GFAP in differentiating cells cultured on laminin after Tg treatment for 6 h and in the absence of Tg for 18 h. Arrows: Dcx-expressing cells; arrowheads: Dcx- and CHOP-expressing cells. Scale bars: 100 µm (a, b). (c) Several neuronal lineage marker proteins, i.e., Dcx, calreticulin, nestin, βIII tubulin, were expressed in differentiating NSCs cultured on laminin, and the expression of the ER stress markers Grp78 and CHOP was induced by Tg. The arrow indicates Dcx. Cells were lysed at 42 h after Tg treatment for 6 h (Tg 6 h + 42 h). D: DMSO. (d) The cells were pretreated with 40 μM zV AD for 30 min, after which Tg (0.23 μM) was added and the cells were incubated for 6 h. After the medium was changed, the cells were incubated with 40 μM zV AD without Tg for 42 h. (e) Semiquantitative RT–PCR at 42 h after treatment with Tg (0.23 μM) for 6 h (upper panels). The mRNA levels of Dcx in cells cultured in medium without Tg were decreased at the indicated times after treatment with 0.23 μM Tg for 6 h (lower panel). The error bars represent the SEMs of 3 independent experiments. (f, g) The mRNA levels of Dcx were not changed after treatment with Tg (0.23 μM) for 6 h in the presence of the IRE1 inhibitor 4μ8C (f) or at 6 h after 0.23 μM of Tg treatment for 6 h in the presence of 40 μM zV AD (g). (b) Semiquantitative RT–PCR analysis of differentiating cells at 6 h after 0.23 μM Tg treatment for 6 h after the knockdown of Dicer using an shRNA and an siRNA (upper panels). Semiquantitative RT–PCR analysis of Dicer (lower left panel, p < 0.05, Student’s t test), shRNA and siRNA for the Control (C) and for Dicer (D). Real-time RT–PCR analysis of Dcx (lower right panel, p < 0.05, Student’s t test). The error bars represent the SEMs.
Discussion

ER stress is activated by several chronic diseases, including obesity, diabetes\(^41\), and depression\(^42\), which may cause dementia later in life\(^41\). However, it is unknown why it takes a long period of time for dementia to develop after body weight increases. In the present study, we demonstrated that long-term obesity induced ER stress, mainly via the ATF4-CHOP axis, in the hippocampal DG, leading to a decrease in the number of processes on Dcx-expressing immature neurons due specifically to the loss of Dcx mRNA stability.

The ATF4-CHOP axis is initiated by eIF2α phosphorylation, which is mediated by four protein kinases, through a process called the ISR, which contributes to the pathogenesis of diseases, including cognitive disorders\(^29\). For example, PERK phosphorylation\(^43\) and eIF2α phosphorylation\(^44\) are observable in AD patients. Consistently, the gene expression of the ISR-related signaling molecule ATF4 is upregulated in the AD brain\(^25\). Since ATF4 directly binds to the promoter of the CHOP (also known as growth arrest and DNA damage-inducible protein, Gadd153) gene\(^45\) and CHOP induction is dependent on ATF4 expression, CHOP is expressed when the ISR is activated. CHOP was detected in the hippocampi of mice with long-term obesity as determined by western blotting (Fig. 2), and immunohistochemistry and confocal microscopy confirmed that CHOP was localized in Dcx-expressing immature neurons of the hippocampal DG (Fig. 3g, h). We observed significantly short processes on Dcx-positive immature neurons in the hippocampal DGs of mice with long-term obesity (Fig. 3). Our experiment showed that the pan-caspase inhibitor zV AD had no effect on the protein expression of Dcx or on Dcx mRNA stability under the ER stress condition (Fig. 4d, g). Further experiments are needed to explore the mechanisms of dendritic loss in Dcx-expressing immature neurons.

ER stress is involved in the pathogeneses of obesity and diabetes\(^46\),\(^47\), which exacerbate cognitive dysfunction in mouse model of AD crossing db/db mice\(^48\). We did not observe the cleavage product of ATF6 or XBP-1s in 20 µg hippocampal lysates even after a longer exposure as determined by western blot (Fig. 2c). Although \(\text{chop}\) expression is regulated by the PERK\(^49\) and ATF6\(^50\) pathways through the binding of ATF4\(^45\) and ATF6\(^51\) to the amino acid-response element (AARE) and ER stress response element (ERSE) of the \(\text{chop}\) promoter, respectively, \(\text{chop}\) mRNA levels were not increased, and \(\text{xbp-1s}\) mRNA was not observed in the hippocampi of aged db/db mice (Fig. 2d), suggesting that ATF6 and Xbp-1 are not involved. The results presented herein demonstrated the potential activation of ER stress/ISR in the immature neurons of mice with long-term obesity. However, the mechanisms of ER stress/ISR activation in immature neurons remain unclear. Reaction oxygen species (ROS) are produced as byproducts of protein oxidation in the ER through ER oxidoreductin-1 (ERO1) and protein disulfide isomerase (PDI)\(^32\). ERO1α transcription is increased by CHOP\(^33\). Overexpression of ERO1β causes ER stress in pancreatic β cells\(^64\). ER stress is also elicited by ROS\(^39\) and by the inactivation of PDI through nitrosylation\(^56\). In the C17.2 NSC line, oxidative stress induced by high-glucose medium activates ER stress and prevents Tuj1 and glial fibrillary acidic protein (GFAP) expression, which is rescued by the superoxide dismutase mimetic tempol and ER stress inhibitor 4-phenylbutyrate\(^37\). Thus, reduction–oxidation (redox) reactions and ER stress stimulate each other\(^29\).

Although we do not know which occurs first in the brains of obese mice, we hypothesize that oxidative stress induces ER stress/ISR in Dcx-expressing immature neurons in vivo. To confirm this hypothesis, further studies are needed; for example, in vivo dynamic nuclear polarization magnetic resonance imaging (DNP-MRI)\(^39\) may allow detection of redox conditions in the brains of young and aged obese mice.

Differentiating NSCs were treated with thapsigargin for 6 h, washed, and cultured in medium without thapsigargin. ER stress was activated in these cells, as indicated by the induction of 78-kDa glucose-regulated protein (Grp78)/immunoglobulin heavy chain-binding protein (BiP) and CHOP expression. The Dcx protein was absent in thapsigargin-treated cell lysates, although the proteins Nestin and βIII Tubulin were present.

**Figure 5.** ER stress increased the miRNA expression in differentiating neurospheres. (a) Differential expression analysis of miRNAs in DMSO-treated cells (n = 3) and thapsigargin-treated cells (n = 3) was performed using edgeR. The y-axis indicates the expression ratio [log2 (fold change)], and the x-axis indicates the average of expression [log (CPM)]. Solid red circle: upregulated miRNA; solid blue circle: downregulated miRNA. FDR < 0.05. (b) Model of the effect of long-term obesity on neural maturation during AHN.
mature neurons of the hippocampal DG (Fig. 5b). Although mRNA stability, resulting in the loss of Dcx in the Dcx-atorning neurospheres under the condition of ER stress (Fig. 5a). These evidences and our results demonstrate that by miRDB72, more experiments are needed to explore the mechanism of mice, which express human to control TAP1 expression in MEFs71. We found that the expression levels of miR-148a-5p, miR-129b-3p, and rhodopsin expression in MEFs70, and Bartoszewski, R., et al. showed that tunicamycin induces diabetes67. It has been suggested that microRNA biogenesis is regulated upon ER stress68. Emde et al. indicated that microRNA biogenesis is inhibited by thapsigargin through the inhibition of DICER activity in the hybrid motor neuron cell line69, in contrast, Behrman et al. demonstrated that thapsigargin induces microRNA-708 to control long-term obesity. Loss of Dcx mRNA by ER stress/ISR during AHN may underlie the memory impairment that occurs later in the lives of obese subjects.

Methods

Western blot analysis and chemicals. Western blotting was performed as described previously32. Mouse tissue and cells were lysed in RIPA [50 mM Tris–HCl (pH 7.5), 150 mM NaCl, 0.1% sodium dodecyl sulfate (SDS), 0.1% sodium deoxycholate, and 1% Nonidet P-40] with Roche cOmplete™ mini tablets (Roche, Mannheim, Germany), 25 μM MG132 (EMD Chemicals, Inc; San Diego, CA), and phosphatase inhibitors, [20 mM β-glycerophosphate (Sigma–Aldrich Co., LLC; St. Louis, MO) and 20 mM sodium orthovanadate (Fujifilm, Osaka, Japan)]. After centrifugation at 13,100 × g, the supernatants were used for western blotting with the following antibodies: anti-CHOP (Thermo Fisher Scientific, Waltham, MA; Cell Signaling Technology, Inc; Danvers, MA), anti-ATF4 (Santa Cruz Biotechnology, Inc; Santa Cruz, CA; Thermo Fisher Scientific), anti-p35 (C-19), anti-GAPDH (Santa Cruz Biotechnology, Inc), anti-a-tubulin (Sigma–Aldrich Co., LLC; St. Louis, MO, anti-Dcx (Abcam, Cambridge, UK; Novus Biologicals, Centennial, CO; Sigma–Aldrich Biotechnology, Inc), anticalreticulin (StressGen Biotechnologies Corp., Victoria, Canada), anti-Nestin (EMD Millipore Corp., Temecula, CA), anti-BiP (Grp78, KDEL) (ENZO Life Sciences International, Inc., Plymouth Meeting, PA). HRP-conjugated anti-rat, anti-mouse, and anti-rabbit IgG (H + L) antibodies (SouthernBiotech; Birmingham, AL) and thapsigargin (Santa Cruz Biotechnology, Inc.) were also used. FBS levels were measured with a FreeStyle Freedom Lite device (Abbott Diabetes Care Inc., Alameda, CA). Insulin levels were measured by a mouse insulin enzyme-linked immunosorbent assay (ELISA) Kit (RTU) (Shibayagi, Gunma, Japan). BDNF levels were measured with a Mature BDNF Rapid™ ELISA Kit (Bisensis Pty Ltd., Thebarton, Australia). All other chemicals were purchased from Wako Pure Chemical Industries, Ltd. (Osaka, Japan), Kanto Chemical Co., Inc., (Tokyo, Japan), and Sigma–Aldrich Co., LLC.

Animals and the HFD. C57BL/6J mice were purchased from Japan SLC, Inc. (Hamamatsu, Japan). APP23 mice, which express human APP751 cDNA and a Swedish double mutation on the C57BL/6 genetic background73, were kindly provided by Dr. M. Staufner (Novartis Pharma Ltd; Basel, Switzerland). Obese and diabetic db/db (Lepr™db/db) mice on the C57BL/6 genetic background were purchased from The Jackson Laboratory (Bar Harbor, ME). The mice were housed in a temperature- and light-controlled room (24 °C; 12:12-light/dark cycle) and fed on AIN93G (standard diet) or HFD-60 (Oriental Yeast Co., Ltd; Tokyo, Japan). All animal studies were approved by the Gifu University Graduate School of Medicine Animal Care and Use Committee and were performed in accordance with the guidelines for experiments on animals provided by the Ministry of Education, Culture, Sports, Science and Technology of Japan. All animal experiments were performed according to the ARRIVE guidelines.

Behavioral tests. The NOL test was performed according to the method described by Roy et al.74. Briefly, each mouse was habituated to a cage without objects for 15 min on day 1. The mice were exposed to three different objects, i.e., conical (diameter x height: 5 x 11.5 cm), cylindrical (6.5 x 10.5 cm), and reagent (5 x 13.2 cm) bottles, for 5 min three times at 2-min intervals on day 2. The mice were placed in their home cages for a retention interval (two minutes). Then, one of the objects was moved to the opposite corner. The behavior of each mouse was monitored using video recording software and an automated tracking system (SMART v3.0 software; Panlab, Barcelona, Spain) and a video camera (HD9-HS350; Panasonic; Osaka, Japan). Three zones (trial1: 3, 4, P; trial4: 3, 4, N) were set at a distance of 4 cm from the object in the far corners of the arena. The exploration
were as follows: 5′-UAC GUA AUU AGA GAG AUC GGCgc-3′. For knockdown of Dicer, we used an siRNA targeting Dicer (sc-108080, Santa Cruz) or a control shRNA (sc-101206, Thermo Fisher Scientific). Differentiating cells were infected with a lentivirus expressing an shRNA targeting Dicer or a control shRNA (sc-108080, Santa Cruz) with 5 µg/mL polybrene at 4 DIV using TransIT-X2 (Takara Bio Inc., Shiga, Japan). The following day, the cells were treated with thapsigargin. The sequences of the siRNAs targeting Dicer were as follows: 5′-GGCGAUCUCUAUUAGCAATT-3′ and 5′-UACGUAUAUGAGGAGCCgc-3′. For immunostaining, the cells were fixed with 4% paraformaldehyde in 0.1 M PB 4°C for 15 min and then incubated in PBS containing 10% normal goat serum and 0.1% Triton X-100 at room temperature for 1 h. The cells were incubated with Alexa Fluor 488-conjugated anti-mouse IgG (H+L) and Alexa Fluor 546-conjugated anti-mouse IgG (H+L) antibodies (Thermo Fisher Scientific). Mouse embryonic stem (ES) cells were grown in DMEM supplemented with 15% normal bovine serum, MEM nonessential amino acids (Thermo Fisher Scientific), 0.1 mM 2-mercaptoethanol (Sigma–Aldrich Co., LLC.), and 1 mM sodium pyruvate (Sigma–Aldrich Co., LLC.). Mouse ES cells were treated with 2 mM DTT for 1 or 30 min, and cells were then lysed by RIPA for western blot analysis.

Small RNA sequencing. Total RNA was purified using a miRNasy mini kit (QIAGEN, Hilden, Germany) from differentiating neurospheres cultured for 6 h after treatment with DMSO or 0.23 µM of thapsigargin for 6 h. Six small RNA-seq libraries (DMSO: n = 3; thapsigargin: n = 3) were generated from the purified total RNA using NEB Next Multiplex Small RNA Library Prep Set for Illumina (Set 1) (New England Biolabs, Ipswich, MA) according to the manufacturer’s instructions. Fifty bp single-end sequencing was performed with NovaSeq

index was calculated as the time spent exploring the object placed at the previous location (P), at a novel location (N), at a zone 3 or at a zone 4 divided by the total time spent in three zones [index P: P/(3 + 4 + P); index N: N/(3 + 4 + N); index 3: 3/(3 + 4 + P or N); index 4: 4/(3 + 4 + P or N)]. When the distance between the nose of the mouse and an object was less than 2 cm or when the mouse sniffed or touched the object with its snout, the mouse was considered to be exploring the object, as previously described. The cages and objects were cleaned with 70% ethanol and 1% acetic acid solution before each trial to eliminate dominant odors.

The MWM test was performed according to a previously described protocol. Briefly, the apparatus was a 100-cm diameter tank containing water at a temperature of approximately 22 °C water, skin milk and a submerged platform. Four acquisition trials from each of the five starting positions were performed each day. The time limit for each trial was 60 s. Mice that did not reach the platform were guided to the platform and left there for 30 s. On day 5, the platform was removed for the probe test. Each trial was recorded and analyzed with the SMART v.3.0 automated tracking system.

Immunostaining. Immunostaining was performed as described previously and analyzed by fluorescence microscopy (BZ-9000, Keyence; Osaka, Japan) and confocal microscopy (LSM710, Carl Zeiss; Göttingen, Germany). Briefly, mice were anesthetized and perfused with phosphate-buffered saline (PBS) and fixed with 4% paraformaldehyde in 0.1 M phosphate buffer (PB). The mouse brains were postfixed for 2 h in the same fixative, which was then replaced with 15% sucrose in 0.1 M PB. The brains were embedded into the optimal critical temperature (OCT) compound (Sakura Finetech USA, Inc., Torrance, CA), and 14-µm-thick sections (coronal plane at a distance of −1.5 to −2.5 mm from bregma: The Mouse Brain in Stereotaxic Coordinates, second edition, Academic Press) were obtained with a cryostat (HM 550, Carl Zeiss). The sections (14 µm) were incubated in PBS supplemented with 10% normal goat serum (Jackson ImmunoResearch Laboratories, Inc., West Grove, PA) and 0.1% Triton X-100 at room temperature for 1 h. The sections were then incubated with an anti-Dcx antibody and an anti-CHOP antibody or an anti-Ki67 antibody (Thermo Fisher Scientific) in PBS containing 1% normal goat serum and 0.1% Triton X-100 at 4 °C for 12 h. For the detection of Ki67, the sections were incubated in 10 mM sodium citrate (pH 6) at 80 °C for 30 min and cooled to room temperature before the antibody was added. On the following days, the sections were incubated with Alexa Fluor 488-conjugated anti-rabbit IgG (H+L) and Alexa Fluor 546-conjugated anti-mouse IgG (H+L) antibodies (Thermo Fisher Scientific) and Hoechst [1 µg/mL bis-benzimide (Sigma–Aldrich Co. LLC.)] to detect fluorescence signals and nuclei. The fluorescence intensity profiles were analyzed with Zen software (Carl Zeiss). The count of Ki67-positive Dcx-positive cells were counted by two people using images obtained by fluorescence microscopy (BZ-9000) and confocal microscopy (LSM710), respectively.

Cell culture and ER stress treatment. Neurons were grown and isolated from the hippocampi of 10-day-old C57BL/6j mice according to a previously described protocol. Neurons were grown in neurobasal medium supplemented with B-27 without vitamin A, 20 ng/mL basic FGF, 20 ng/mL EGF, GlutaMAX, and gentamicin (Thermo Fisher Scientific). For differentiation, the cells were plated on dishes coated with natural mouse laminin (Thermo Fisher Scientific); the next day, the medium was replaced with medium A, which was comprised of Dulbecco’s modified Eagle’s medium and Ham’s F-12 (DMEM/F12, Wako Pure Chemical Industries, Ltd.), MACS® NeuroBrew®-21 (Miltenyi Biotec, Bergisch Gladbach, Germany), 0.5 × N-2 supplement (Thermo Fisher Scientific), 20 ng/mL basic FGF, and gentamicin. On the following days, the medium A was changed in the morning and replaced with medium B, which consisted of DMEM/F12, MACS® NeuroBrew®-21, 0.5 × N-2 supplement, and gentamicin, in the afternoon. Then, the medium was changed every two days. The cells were maintained at 37 °C in an atmosphere containing 5% CO₂. Thapsigargin was added to the medium at a final concentration of 0.23–0.69 µM, after which the cells were incubated in 6 h and washed with medium. The medium was replaced with new medium, and the cells were incubated for the indicated amounts of time. Knockdown of Dicer was induced by an shRNA-expressing lentiviral vector (Dicer, Santa Cruz) and an siRNA (Dicer, Thermo Fisher Scientific). Differentiating cells were infected with a lentivirus expressing an shRNA targeting Dicer or a control siRNA (sc-108080, Santa Cruz) with 5 µg/mL polybrene at 1 DIV and then transfected with an siRNA targeting Dicer or a control siRNAs at 4 DIV using TransIT-X2 (Takara Bio Inc., Shiga, Japan). The following day, the cells were treated with thapsigargin. The sequences of the siRNAs targeting Dicer were as follows: 5′-GCACUCAUCAUCAGGAATT-3′ and 5′-UACGUAUAUGAGGAGCGgc-3′. For immunostaining, the cells were fixed with 4% paraformaldehyde in 0.1 M PB 4°C for 15 min and then incubated in PBS containing 10% normal goat serum and 0.3% Triton X-100 at room temperature for 1 h. The cells were incubated with an anti-Dcx antibody, an anti-CHOP antibody and an anti-GFAP antibody (Thermo Fisher Scientific) in PBS containing 1% normal goat serum and 0.1% Triton X-100 at 4 °C for 12 h. The cells were then incubated with Alexa Fluor 488-conjugated anti-rabbit IgG (H+L) and Alexa Fluor 546-conjugated anti-mouse IgG (H+L) antibodies (Thermo Fisher Scientific).

Small RNA sequencing. Total RNA was purified using a miRNasy mini kit (QIAGEN, Hilden, Germany) from differentiating neurospheres cultured for 6 h after treatment with DMSO or 0.23 µM of thapsigargin for 6 h. Six small RNA-seq libraries (DMSO: n = 3; thapsigargin: n = 3) were generated from the purified total RNA using NEB Next Multiplex Small RNA Library Prep Set for Illumina (Set 1) (New England Biolabs, Ipswich, MA) according to the manufacturer’s instructions. Fifty bp single-end sequencing was performed with NovaSeq.
Semiquantitative and quantitative RT–PCR. RNA was isolated from cells with TRIzol (Thermo Fisher Scientific) as described previously. Reverse transcription was performed using M-MLV reverse transcriptase (Thermo Fisher Scientific) with random primers (Toyobo Co., Ltd., Osaka, Japan). RT–PCR analysis was performed using an S1000 Thermal Cycler (Bio–Rad, Hercules, CA) with Takara Ex Taq (Takara, Shiga, Japan). Semiquantitative polymerase chain reaction (PCR) was performed for 25 cycles. Quantitative RT–PCR was performed using a TP870 Thermal Cycler Dice (Takara) with Thunderbird SYBR qPCR Mix (TOYOBO CO., LTD. Osaka, Japan). The PCR primer pairs were as follows: mouse Dcx: 5′-GCTACATTITACCATGTGAGGATCCAG3′ and 5′-TCATCATCAAGAGAATCTGAGACGGATGC3′; mouse Nestin: 5′-GAAGTCAGACTGGCTCAAGTCC3′ and 5′-GGAGGACACCTAGTAACTAGG3′; mouse p35: 5′-CTGCGACCCCATCCCTCACATC3′ and 5′-GAACACCTTATGTCATGAGGCTTTG3′; mouse Xbp-1: 5′-GAAGTCGCGAAAAAGATCATGAGCTC3′ and 5′-GGCTCTGTGTTGAGAACCAGGAG3′; mouse Dicer: 5′-AGACACACCTGTCTATGGCAAC3′ and 5′-CAATCCCTCGTGCACCTTTGAC3′; and mouse β–Actin: 5′-CTCAAAGGCAACCGTGAAAG3′ and 5′-CAGCGACGATTTTCTCTCA3′.

Statistics. Statistical analyses were performed using SPSS Statistics 27 (IBM, Armonk, NY) and Excel (Microsoft, Redmond, WA). Statistical significance (p<0.05) was determined using Student’s t test (two-tailed) or one-way ANOVA followed by the Bonferroni post-hoc test. The statistical details are summarized in Supplementary Table 2.

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Author contributions
K.N. and T.N. contributed to the experimental design and data interpretation, and K.N., S.I., M.U., and T.N. contributed to the data analysis. All authors critically read and approved the manuscript.

Competing interests
The authors declare no competing interests.

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