

Expression of zinc finger protein ZPR1 mRNA in brain is up-regulated in mice fed a high-fat diet

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Received August 16, 2005; Accepted September 23, 2005

Abstract. A differential display was performed to analyze differential gene expression in the brain of mice in association with dietary high beef tallow. Consumption of a high beef tallow diet up-regulated the expression of zinc finger protein ZPR1 mRNA in the brain. Expression of ZPR1 mRNA in the cerebellum and hippocampus was elevated in response to the high beef tallow diet. The increased ZPR1 expression in the neuronal, Neuro-2A cells, caused a significant increase in H₂O₂-induced cell death. These results suggest that a high beef tallow diet up-regulates ZPR1 mRNA expression in the brain and might increase the vulnerability to oxidative stress.

Introduction

Fats constitute about 40% of the dietary energy intake in most industrialized Western societies. High intake of animal fat is a risk factor for the development of not only coronary heart disease, diabetes, and cancer (1,2), but also brain diseases including Alzheimer's disease (AD) and Parkinson's disease (PD) (3-7). Morris *et al* reported that intake of saturated and trans-unsaturated fatty acids was positively associated with the risk of AD, whereas intake of ω -6, ω -3 polyunsaturated, and monosaturated fatty acids was inversely associated with this risk (3,4). Also, several studies have suggested positive correlations of PD risk with the consumption of total fat, saturated fatty acids, oleic acids, and cholesterol (5-7). In addition, some epidemiological studies have revealed that high intake of total fat, saturated fatty acids, and cholesterol is associated with an increased risk of dementia (8,9).

Animal studies have demonstrated that cognitive and leaning performance is affected by simple alterations in dietary fatty acid composition. Greenwood *et al* demonstrated that consumption of a diet high in saturated fatty acids impairs cognitive functions in rats fed diets containing 40% of energy from various fat blends for three months (10,11). Molteni *et al* demonstrated that consumption of a diet high in saturated fatty acids for two months impairs leaning performance in rats through down-regulation of several genes, including hippocampal brain-derived neurotrophic factor (BDNF), which is a crucial mediator of neuronal viability and function involved in learning and memory (12). Therefore, elevation of risk for brain disease and cognitive decline induced by a diet high in saturated fatty acids may be mediated by modulation of gene expression in brain. Further searches for dietary fat-responsive genes involving neuronal viability and function are required to elucidate the mechanisms of this increased risk of brain disease and cognitive decline.

The aim of the present study was to identify neuronal genes regulated in response to consumption of a diet containing beef tallow rich in saturated fatty acids using a differential display method. Here, we show that short-term feeding of a high beef tallow diet up-regulated the expression of zinc finger protein ZPR1 (ZPR1) mRNA in brains of mice. ZPR1 is a zinc finger protein that acts as a signaling factor relaying proliferative growth signals from the cell surface to the nucleus (13,14). Our study further demonstrated that higher expression of ZPR1 in the neuronal, Neuro-2A cells, increased H₂O₂-induced cell death, suggesting that the higher consumption of a saturated fat diet enhances neuronal vulnerability to oxidative stress through elevation of ZPR1 gene expression.

Materials and methods

Materials. Purified corn oil and beef tallow were purchased from Nacalai Tesque, Inc. (Kyoto, Japan). Restriction endonucleases and DNA-modifying enzymes were products of Takara Bio (Kyoto, Japan). Dulbecco's modified Eagle's medium (DMEM), fetal calf serum (FCS), and G418 were purchased from Invitrogen. Horseradish peroxidase-conjugated anti-rabbit IgG antibody was purchased from Oncogene.

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Abbreviations: AD, Alzheimer's disease; PD, Parkinson's disease; ZPR1, zinc finger protein ZPR1; EGF, epidermal growth factor; GST, glutathione S-transferase; EF-1 α , eukaryotic translation elongation factor 1 α ; SMN, survival motor neuron protein

Key words: high fat diet, brain, ZPR1

Animals and diets. Male CD-1 (ICR) mice (6-weeks-old, Charles River, Japan) were housed in groups of three animals

in a metal cage in a room with controlled temperature (24°C) and a 12-h light:dark cycle (lights on 08:00-20:00 h). Animals were given free access to diet and deionized water. The mice were maintained according to the Guide for the Care and Use of Laboratory Animals established by Hiroshima University. After being fed a commercial stock diet (MF, Oriental Yeast, Tokyo, Japan) for 1 week, the mice (average 28 g) were divided into groups of 9 mice. The composition of the experimental diets is presented in Table I. The feeding experimental period was 28 days.

Differential display. Total RNA was isolated using an RNeasy lipid tissue kit (Qiagen), and differential display was performed as previously described (15). Briefly, 5 µg of total RNA was converted to cDNA using a First strand reaction cDNA synthesis kit (Amersham Bioscience). Subsequently, each cDNA was amplified by PCR with 100 different arbitrary primers. After separation by polyacrylamide gel electrophoresis (PAGE), cDNA bands that appeared to be up-regulated in the HB group were recovered and reamplified with the corresponding primer. The cDNAs were sequenced by the dideoxy chain termination method using an Applied Biosystems model 373A automated sequencer and a dye terminator cycle sequencing kit (Amersham Bioscience). The complete cDNA of ZPR1 was cloned by PCR amplification with a mouse brain cDNA library (Clontech) as a template and primers 5'-GAAGCCGCCGCGGGAGCAACGCGTG-3' and 5'-TGCAGTGTGAGTGGCACTGTAGCTG-3'.

Northern blot analysis and in situ hybridization. Total RNA (10 µg) was separated by electrophoresis in a 1% agarose/2.2 M formaldehyde denaturing gel, and were then transferred to a Hybond N⁺ nylon membrane (Amersham Bioscience). ³²P-labeled cDNA fragments encoding mouse ZPR1 and human β-actin were used for Northern hybridization as a probe. Hybridization was performed in 6X sodium saline citrate (SSC), 0.5% SDS, 5X Denhardt's solution, 100 µg/ml denatured salmon sperm DNA at 65°C for 16 h. The membrane was washed with 2X SSC/0.1% SDS at 25°C, and 0.2X SSC/0.1% SDS at 65°C. The membrane was exposed to a Fuji imaging plate (FUJI photo film, Japan) for an appropriate time. The radioactivity in each band was quantified using an image analyzer and normalized to that of β-actin mRNA. *In situ* hybridization was performed using brains of male ICR mice and digoxigenin-labeled probes as described previously (16). Brains of mice were rapidly fixed with paraformaldehyde, and 15-µm-thick sections were prepared on a cryostat. Digoxigenin-labeled cDNA probes (antisense and sense) were made by *in vitro* transcription using cDNAs subcloned into pGEM-T vector as templates in the presence of digoxigenin-labeled dUTP (Roche) according to the manufacturer's instructions.

Preparation of recombinant ZPR1 protein and anti-ZPR1 antibody. The open reading frame of mouse ZPR1 was subcloned in-frame into the glutathione S-transferase (GST) expression vector pGEX-6P-3 (Amersham Bioscience), generating pGEX-ZPR1. The pGEX-ZPR1 was introduced into the bacterial strain JM109 (Toyobo, Japan). Over-expression of GST-fused ZPR1 protein was performed as

Table I. Composition of experimental diets.

Ingredients (g/kg)	LC	LB	HB	UHB
Corn oil	50	0	0	0
Beef tallow	0	50	200	350
Corn starch	452	452	302	152
Casein	200	200	200	200
L-Cystine	3	3	3	3
Cellulose powder	50	50	50	50
Vitamin mixture ^a	10	10	10	10
Salt mixture ^b	35	35	35	35
Sucrose	200	200	200	200

^aAIN-93, ^bAIN-93G.

described previously (16). The suspended cells were sonicated in an ice-cold solubilization buffer (50 mM Tris-HCl, pH 8.0, and 1 mM EDTA), and the homogenates were cleared by centrifugation at 15,000 x g for 15 min at 4°C. The supernatant was then purified on a column of Glutathione Sepharose 4 Fast Flow (Amersham Bioscience) according to the manufacturer's protocol. Polyclonal antibody raised against ZPR1 was obtained by injecting purified GST-fused ZPR1 into rabbits with Freund's complete adjuvant (DIFCO).

Transfection of ZPR1 into Neuro-2A cells and H₂O₂-induced cell death in stable transfectants. The open reading frame of ZPR1 was subcloned in-frame into pEF/myc/cyto (Invitrogen), adding a C-terminal c-myc tag to ZPR1 (pEF/ZPR1-myc). Neuro-2A cells were cultured in DMEM with 10% fetal calf serum, 100 units/ml penicillin, and 100 µg/ml streptomycin at 37°C/5% CO₂, and were transfected with the pEF/ZPR1-myc (ZPR-myc) or the empty pEF/myc/cyto vector (mock) using Lipofectamine 2000 (Invitrogen). For establishment of stable transfectants, Neuro-2A cells were treated with 500 µg/ml G418 for 14 days. G418-resistant colonies were identified and re-seeded in DMEM with 10% fetal calf serum and 200 µg/ml G418 at 37°C/5% CO₂. After the stable transfectants expressing ZPR1-myc protein and mock cells were seeded in 24-well dishes at 2.5x10⁴ cells/cm², the cells were exposed to 200 µM H₂O₂ for 6 h. The stable transfectants were washed in PBS and harvested by scraping in 400 µl of PBS. After centrifugation for 5 min at 100 x g, the pellets were resuspended in 100 µl of PBS, and subjected to cell counting with a hemocytometer.

Western blot analysis. Neuro-2A cells were washed with ice-cold PBS, and scraped in an ice-cold RIPA buffer (10 mM Tris-HCl, pH 7.4, 1% NP-40, 0.1% sodium deoxycholate, 0.1% SDS, 0.15 M NaCl, 1 mM EDTA). The homogenates were centrifuged at 10,000 x g for 10 min. Protein concentration of the supernatant was determined by the Bio-Rad protein assay kit (Bio-Rad) with BSA as standard. The supernatant (10 µg, protein equivalents) was subjected to

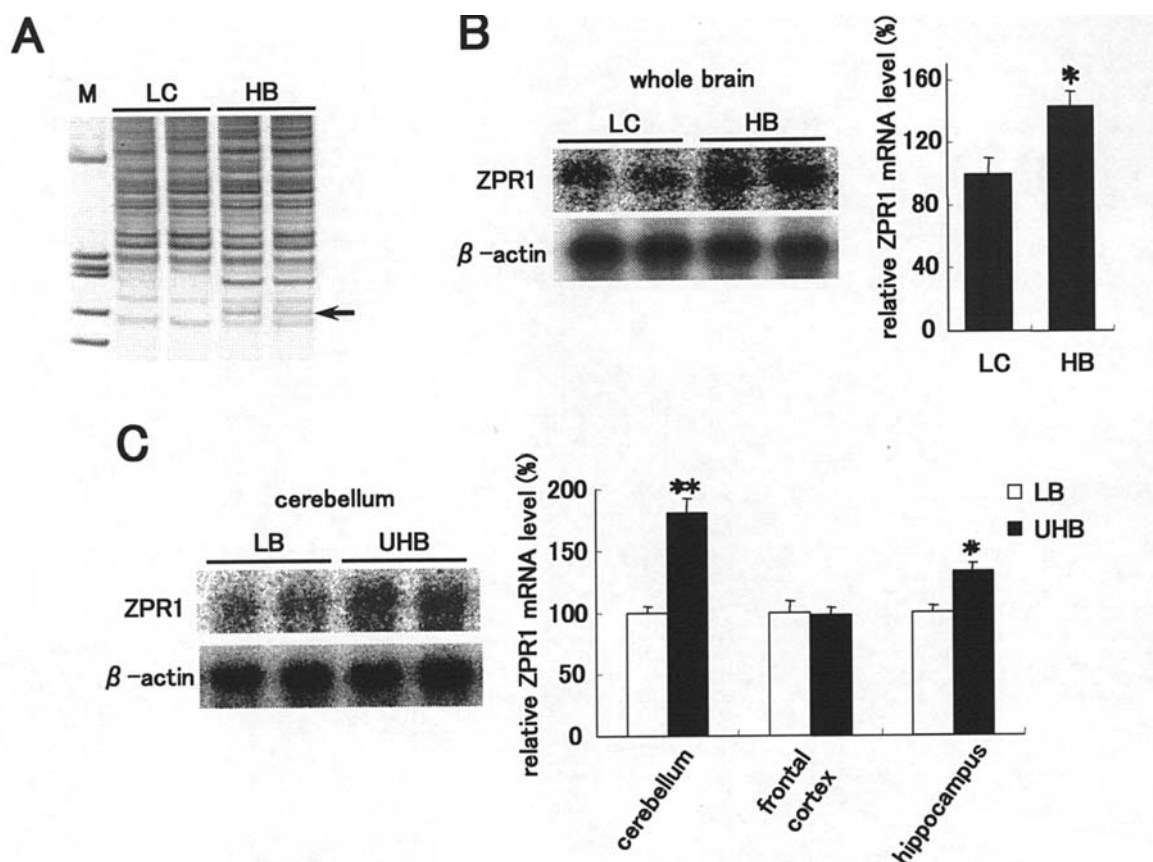


Figure 1. Differential expression of ZPR1 mRNA in mouse brains. A, Differential display PCR of cDNA from brains of mice fed a low corn oil diet (LC) or high beef tallow diet (HB). B, Northern blot analysis for ZPR1 mRNA in the whole brains of mice fed LC or HB. Total RNAs (10 μ g) were subjected to Northern blot analysis. In the autoradiograms, each lane represents a sample from an individual mouse. The radioactivity in each band was quantified using an image analyzer and normalized to that of β -actin mRNA. Each value is expressed relative to the mRNA level of the LC group and represents the mean \pm SE (n=3, *P<0.05 vs. LC group). C, Northern blot analysis for ZPR1 mRNA in the cerebellum, frontal cortex or hippocampus of mice fed a low beef tallow diet (LB) or ultra high beef tallow diet (UHB). In autoradiograms, each lane represents a sample from the cerebellum of an individual mouse. The radioactivity in each band was quantified using an image analyzer and normalized to that of β -actin mRNA. Each value is expressed relative to the mRNA level of the LB group and represents the mean \pm SE (n=3, *P<0.05, **P<0.01 vs. LB group).

SDS-PAGE and transferred to an Immobilon P filter (Millipore). The filter was blocked for 18 h at 4°C by soaking in 4% nonfat dried milk in PBS and was incubated for 18 h at 4°C with anti-ZPR1 anti-body (diluted 1:1000). Signals were detected using horseradish peroxidase-conjugated anti-rabbit IgG and the enhanced chemiluminescence system (Amersham Bioscience).

Data analysis. In all experiments, means \pm SEM are reported. Statistical comparison for the experiment of cell survival among clones transfected with ZPR1 or empty vector was determined using one-way analysis of variance (one-way ANOVA) and Duncan's multiple-range test (17). Statistical comparison for other experiments was analyzed by Student's t-test.

Results

Isolation of genes up-regulated in response to a high beef tallow diet. After 4 weeks of the diet, no significant differences were found in body weight, food intake, and the weights of epididymal and perirenal adipose tissues between the LC and HB groups (body weight: LC, 40.9 \pm 1.3 g; HB, 41.1 \pm 1.8 g; food intake: LC, 5.0 \pm 0.8 g/day; HB, 4.3 \pm 0.7 g/day; adipose tissue weight: LC, 2.02 \pm 0.14 g; HB, 2.06 \pm 0.26 g).

To identify genes that were differentially expressed between the LC and HB group, we performed the differential display procedure. The whole brain cDNAs were prepared from two mice in each of these dietary groups, and were subjected to differential display PCR reactions in the presence of different sets of primers with arbitrary sequences. We focused on a cDNA fragment (S6) that was found to be amplified to a greater degree from the whole brain cDNA of the HB group (Fig. 1A). The S6 fragment was reamplified and subcloned into a TA cloning vector. As a result of nucleotide sequencing of the clone, the S6 fragment was identified as a part of mouse zinc finger protein ZPR1 cDNA. To confirm the differential expression of the ZPR1 gene, total RNA from individual mice of the LC and HB groups was subjected to Northern blot analysis (Fig. 1B). The ZPR1 mRNA level was significantly up-regulated in the HB group compared with the LC group (with a 42% increase, P<0.05, Fig. 1B).

Beef tallow-responsiveness of expression of ZPR1 mRNA in distinct brain regions. In order to analyze the effects of beef tallow intake on ZPR1 mRNA expression in the brain, Northern blot analysis for expression of ZPR1 mRNA was further employed in cerebellum, frontal cortex, and hippocampus from mice fed either LB or UHB. Fig. 1C shows that

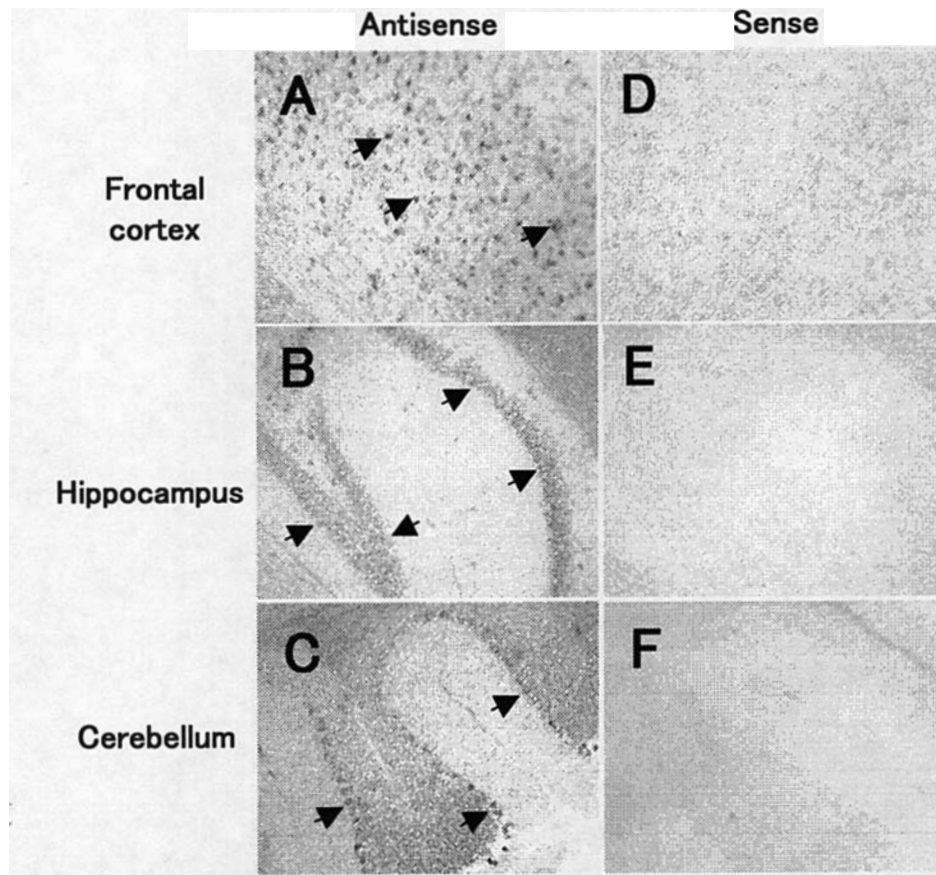


Figure 2. Distribution of ZPR1 mRNA expression in the normal mouse brain. *In situ* localization of ZPR1 mRNA in mouse brain. Frozen sections of normal mouse brain were hybridized with digoxigenin-labeled antisense or sense cDNA probe complementary to ZPR1 mRNA as described under Materials and methods. For detection of hybridized cDNA probes, anti-digoxigenin antibody conjugated to alkaline phosphatase was used, and color was developed by incubation with 4-nitro blue tetrazolium chloride and X-phosphate solution. Expression of ZPR1 mRNA was observed in the frontal cortex, cerebellum, and hippocampus regions (arrows). In particular, strong signals were detected in most Purkinje cells and in the cornus ammonis and dentate gyrus of the hippocampus.

the level of ZPR1 mRNA was significantly up-regulated in the cerebellum (with an 81% increase, $P < 0.01$) and hippocampus (with a 31% increase, $P < 0.05$) regions of the UHB group compared with those of the LB group. No difference of the expression of ZPR1 mRNA was seen in the frontal cortex region.

Expression of ZPR1 mRNA in mouse brain. To identify the cell types in which ZPR1 mRNA is expressed, we performed *in situ* hybridization with ZPR1 cDNA probe and frozen sections from mouse brain. As shown in Fig. 2, ZPR1 mRNA was detected in the frontal cortex (Fig. 2A), cornus ammonis and dentate gyrus of the hippocampus (Fig. 2B), and in Purkinje cells of the cerebellum (Fig. 2C), indicating that neuronal ZPR1 mRNA is mainly expressed in neurons.

Preparation of anti-ZPR1 antibody and stable transfectants of Neuro-2A cells that overexpress mouse ZPR1. A polyclonal antibody against a ZPR1 protein was prepared using a bacterially synthesized GST-fused protein. The specificity of the anti-ZPR1 antibody was tested by Western blot analysis of total proteins from COS-7 cells transiently transfected with an expression vector encoding ZPR1-myc. Although no signal was observed in the mock-transfected COS-7 cells, a specific band was detected in the cells transfected with the

expression vector encoding ZPR1-myc using anti-ZPR1 antibody (data not shown). The Neuro-2A cells are extensively used as neuronal models *in vitro*. In addition, the endogenous expression of ZPR1 mRNA was observed in Neuro-2A cells (data not shown). To investigate whether the up-regulation of ZPR1 mRNA expression influences cell survival in the neuronal cell line, we established stably transfected clones of Neuro-2A cells that overexpress mouse ZPR1. As shown in Fig. 3B, elevation of ZPR1 protein was observed in stably transfected clones, Z3 and Z14, using the anti-ZPR1 polyclonal antibody, but not in mock clones M1 and M2. The expression of ZPR1 mRNA was also up-regulated in the ZPR1-transfected clone Z3 compared with the mock clone M1 (Fig. 3A).

Effect of increased ZPR1 expression on H_2O_2 -induced cell death in Neuro-2A cells. Fig. 3C shows cell survival of the stable transfectants of Neuro-2A exposed to $200 \mu M H_2O_2$ for 6 h. Exposure to H_2O_2 resulted in the reduction of cell survival, with a 72 and a 65% decrease in the Z3 and Z14 clones, respectively, whereas the exposure resulted in only a 48 and a 45% decrease in the M1 and M2 clones, respectively. There were no significant differences in cell survival of M1, M2, Z3, and Z14 clones without H_2O_2 for 6 h (data not shown). These

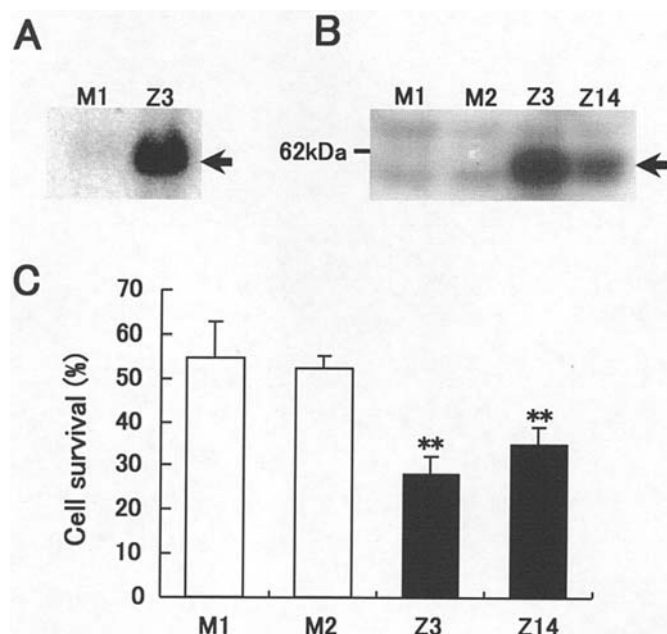


Figure 3. Survival of stable transfectants exposed to H₂O₂. A, ZPR1 mRNA was up-regulated in clone ZPR1-3 (Z3) cells compared with clone mock1 (M1) cells. Total RNA (10 μ g) from Z3 and M1 cells were subjected to Northern blot analysis. B, ZPR1 protein was elevated in clone ZPR1-3 (Z3) and ZPR1-14 (Z14) cells compared with clone mock1 (M1) and mock2 (M2) cells. Whole-cell extracts prepared from Z3, Z14, M1, and M2 cells were subjected to SDS-PAGE, followed by Western blot analysis with anti-ZPR1 antibody. C, These transfectants were incubated with 200 μ M H₂O₂ for 6 h, followed by measurement of H₂O₂-induced cell death. Values are the mean \pm SE (n=5, **P<0.01 vs. M1).

results suggest that up-regulation of ZPR1 expression increases their vulnerability to oxidative stress.

Discussion

There is increasing evidence that higher consumption of animal fat is involved in the pathogenesis of brain diseases and the impairment of neuronal function. In particular, a higher intake of saturated fatty acids has been shown to elevate the risk of AD and PD and to impair cognitive function. However, the molecular mechanisms by which dietary fat influences the brain are poorly understood. In the present study, a differential display method was performed to isolate genes in the mouse brains in response to dietary high beef tallow. Herein it is demonstrated that ZPR1 mRNA expression is up-regulated in the brains of mice fed high beef tallow diets.

ZPR1 is an essential protein whose amino acid sequence is highly conserved from yeast to mammals (18). ZPR1 has two zinc finger motifs which bind to epidermal growth factor (EGFR) and platelet-derived growth factor receptor (PDGFR) (13). Previous reports have shown that treatment with mitogens induced the release of ZPR1 from the growth factor receptors and translocation to the nucleus (14). Therefore, ZPR1 is supposed to act as a signaling factor relaying proliferative growth signals from the cell-surface to the nucleus. In addition, ZPR1 is shown to bind to eukaryotic polypeptide elongation factor 1 α (EF-1 α) and survival motor neuron protein (SMN) in cytoplasm (18,19). Previous studies demonstrated that the complex with ZPR1, EF-1 α and SMN translocates to the

nucleus after stimulation of serum. EF-1 α is known to be involved in the transfer of aminoacyl-tRNA to the ribosome, cytoskeletal organization, apoptosis, and nuclear processes, such as RNA synthesis and mitosis (20). Gangwani *et al* reported that disruption of the binding of ZPR1 to EF-1 α results in an accumulation of cells in the G2/M phase of cell cycle and defective growth (18). While, SMN protein implicated in several important physiological pathways, including nucleo-cytoplasmic RNA transportation, uridine-rich small nuclear ribonucleoprotein assembly, RNA transcription, and apoptosis (21-24). Thus, up-regulation of ZPR1 expression might influence the cell cycle, apoptosis and RNA metabolism in neurons.

Neuronal cell death is increasingly recognized as a major contributor to neuropathology such as AD and PD underlying a variety of degenerative diseases of the brain. The cell death of neurons in these disorders is thought to be caused by toxic insults such as abnormal protein aggregates and free-radical generation. The present study demonstrated that up-regulation of ZPR1 expression in Neuro-2A enhanced the H₂O₂-induced cell death, suggesting that higher expression of ZPR1 increases the vulnerability to oxidative stress in neurons. Several studies have demonstrated that oxidative stress is a major risk factor for the initiation and progression of sporadic PD and AD (25,26). Because intake of high fat diets, obesity, and diabetes can elevate oxidative stress in plasma and brain (27,28), these observations imply that high animal-fat diet results in cognitive decline and neuronal disorders by elevating oxidative stress in combination with an increase in vulnerability to oxidative stress. Although this study opens important clues to the elucidation of the impact of dietary fat on the development of brain diseases, elucidation of the physiologic function of ZPR1 in neurons will be required for understanding the molecular basis of the effects of high fat diets on the brain functions.

References

- Mann JJ: Diet and risk of coronary heart disease and type 2 diabetes. *Lancet* 360: 783-789, 2002.
- Liu Z, Uesaka T, Watanabe H and Kato N: High fat diet enhances colonic cell proliferation and carcinogenesis in rats by elevating serum leptin. *Int J Oncol* 19: 1009-1014, 2001.
- Morris MC, Evans DA, Bienias JL, Tangney CC, Bennett DA, Aggarwal N, Schneider J and Wilson RS: Dietary fats and the risk of incident Alzheimer disease. *Arch Neurol* 60: 194-200, 2003.
- Morris MC, Evans DA, Bienias JL, Tangney CC, Bennett DA, Wilson RS, Aggarwal N and Schneider J: Consumption of fish and n-3 fatty acids and risk of incident Alzheimer disease. *Arch Neurol* 60: 940-946, 2003.
- Logroscino G, Marder K, Cote L, Tang MX, Shea S and Mayeux R: Dietary lipids and antioxidants in Parkinson's disease: a population-based, case-control study. *Ann Neurol* 39: 89-94, 1996.
- Johnson CC, Gorell JM, Rybicki BA, Sanders K and Peterson EL: Adult nutrient intake as a risk factor for Parkinson's disease. *Int J Epidemiol* 28: 1102-1109, 1999.
- Anderson C, Checkoway H, Franklin GM, Beresford S, Smith-Weller T and Swanson PD: Dietary factors in Parkinson's disease: the role of food groups and specific foods. *Mov Disord* 14: 21-27, 1999.
- Kalmijn S: Fatty acid intake and the risk of dementia and cognitive decline: a review of clinical and epidemiological studies. *J Nutr Health Aging* 4: 202-207, 2000.
- Heude B, Ducimetiere P and Berr C: Cognitive decline and fatty acid composition of erythrocyte membranes - the EVA Study. *Am J Clin Nutr* 77: 803-808, 2003.

10. Greenwood CE and Winocur G: Learning and memory impairment in rats fed a high saturated fat diet. *Behav Neural Biol* 53: 74-87, 1990.
11. Greenwood CE and Winocur G: Cognitive impairment in rats fed high-fat diets: a specific effect of saturated fatty-acid intake. *Behav Neurosci* 110: 451-459, 1996.
12. Molteni R, Barnard RJ, Ying Z, Roberts CK and Gomez-Pinilla F: A high-fat, refined sugar diet reduces hippocampal brain-derived neurotrophic factor, neuronal plasticity, and learning. *Neuroscience* 11: 803-814, 2002.
13. Galcheva-Gargova Z, Konstantinov KN, Wu IH, Klier FG, Barrett T and Davis RJ: Binding of Zinc finger protein ZPR1 to the epidermal growth factor receptor. *Science* 272: 1797-1802, 1996.
14. Galcheva-Gargova Z, Gangwani L, Konstantinov KN, Mikrut M, Theroux SJ, Enoch T, and Davis RJ: The cytoplasmic zinc finger protein ZPR1 accumulates in the nucleolus of proliferating cells. *Mol Biol Cell* 9: 2963-2971, 1998.
15. Yanaka N, Imai Y, Kawai E, Akatsuka H, Wakimoto K, Nogusa Y, Kato N, Chiba H, Kotani E, Omori K and Sakurai N: Novel membrane protein containing glycerophosphodiester phosphodiesterase motif is transiently expressed during osteoblast differentiation. *J Biol Chem* 278: 43595-43602, 2003.
16. Yuasa K, Omori K and Yanaka N: Binding and phosphorylation of a novel germ cell-specific cGMP-dependent protein kinase-anchoring protein by cGMP-dependent protein kinase Ialpha. *J Biol Chem* 275: 4897-4905, 2000.
17. Duncan DB: Multiple range test for correlated and heteroscedastic means. *Biometrics* 13: 164-176, 1957.
18. Gangwani L, Mikrut M, Galcheva-Gargova Z and Davis RJ: Interaction of ZPR1 with translation elongation factor-1 α in proliferating cells. *J Cell Biol* 143: 1471-1484, 1998.
19. Gangwani L, Mikrut M, Theroux S, Sharma M and Davis RJ: Spinal muscular atrophy disrupts the interaction of ZPR1 with SMN protein. *Nat Cell Biol* 3: 376-383, 2001.
20. Ejiri S: Moonlighting functions of polypeptide elongation factor 1: from actin bundling to zinc finger protein ZPR1-associated nuclear localization. *Biosci Biotech Biochem* 66: 1-21, 2002.
21. Schmalbruch H and Haase G: Spinal muscular atrophy: present state. *Brain Pathol* 11: 231-247, 2001.
22. Paushkin S, Gubitz AK, Massenet S and Dreyfuss G: The SMN complex, an assemblyosome of ribonucleoproteins. *Curr Opin Cell Biol* 14: 305-312, 2002.
23. Pellizzoni L, Charroux B, Rappaport J, Mann M and Dreyfuss G: A functional interaction between the survival motor neuron complex and RNA polymerase II. *J Cell Biol* 152: 75-85, 2001.
24. Gisemann T, Rathke-Hartlieb S, Rothkegel M, Bartsch JW, Buchmeier S, Jockusch BM and Jockusch H: A role for polyproline motifs in the spinal muscular atrophy protein SMN. Profilins bind to and colocalize with smn in nuclear gems. *J Biol Chem* 274: 37908-37914, 1999.
25. Zhu X, Raina AK, Lee HG, Casadesus G, Smith MA and Perry G: Oxidative stress signalling in Alzheimer's disease. *Brain Res* 1000: 32-39, 2004.
26. Schapira AH: Disease modification in Parkinson's disease. *Lancet Neurol* 6: 362-368, 2004.
27. Opera EC: Role of oxidative stress in the etiology of type 2 diabetes and the effect of antioxidant supplementation on glycemic control. *J Invest Med* 108: 175-180, 2004.
28. Beltowski J, Wojcik G, Gorny D and Marciniak A: The effects of dietary-induced obesity on lipid peroxidation, antioxidant enzymes and total plasma antioxidant capacity. *J Physiol Pharmacol* 51: 883-896, 2000.