NEUROBIOLOGY

Ganglioside Synthase Knockout Reduces Prion Disease Incubation Time in Mouse Models

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Localization of the abnormal and normal isoforms of prion proteins to detergent-resistant membrane microdomains, lipid rafts, is important for the conformational conversion. Lipid rafts are enriched in sialic acid-containing glycosphingolipids (namely, gangliosides). Alteration in the ganglioside composition of lipid rafts can affect the localization of lipid raft-associated proteins. To investigate the role of gangliosides in the pathogenesis of prion diseases, we performed intracerebral transmission study of a scrapie prion strain Chandler and a Gerstmann-Sträussler-Scheinker syndrome prion strain Fukuoka-1 using various knockout mouse strains ablated with ganglioside synthase gene (ie, GD2/GM2 synthase, GD3 synthase, or GM3 synthase). After challenge with the Chandler strain, GD2/GM2 synthase knockout mice showed 20% reduction of incubation time, reduced prion protein deposition in the brain with attenuated glial reactions, and reduced localization of prion proteins to lipid rafts. These results raise the possibility that the gangliosides may have an important role in prion disease pathogenesis by affecting the localization of prion proteins to lipid rafts. (*Am J Pathol* 2019; 189: 677–686; <https://doi.org/10.1016/j.ajpath.2018.11.009>)

Prion diseases are fatal transmissible neurodegenerative diseases, including Creutzfeldt-Jakob disease, Gerstmann-Sträussler-Scheinker syndrome, and fatal familial insomnia in humans, or bovine spongiform encephalopathy, chronic wasting disease, and scrapie in animals. Accumulation of an abnormal protease-resistant isoform of prion protein (PrP^{Sc}), which is generated by conformational conversion of the normal cellular isoform (PrP^C), is one of the pathognomonic features of prion diseases.¹

Membrane microdomains where PrP^C and PrP^{Sc} are localized are lipid rafts.^{2–6} Lipid rafts are enriched in cholesterol, sphingolipids, and glycosylphosphatidylinositol (GPI)-anchored proteins, including PrP, and show partial resistance against mild detergent treatment.^{7,8} Therefore, lipid rafts buoy toward low-density fractions in density gradient ultracentrifugation after mild detergent treatment (namely, flotation assay). Lipid rafts have been reported to

serve many microorganisms for infection or serve the clustering of receptors or signaling molecules to facilitate signal transduction.^{8,9} In prion disease pathogenesis, the localization of PrP^C and PrP^{Sc} to lipid rafts has been reported to be important for the conformational conversion.³

Gangliosides are sialic acid-containing glycosphingolipids highly expressed in the brain. Gangliosides are synthesized in the Golgi apparatus by specific glycosyltransferases, the expression of which controls the steady-state levels of gangliosides.¹⁰ Gangliosides are characterized by the number and positions of sialic acids that define their

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classification into A, M, D, T, Q, and P (zero to five sialic acids) and a, b, and c series (one, two, or three sialic acids on internal galactose residue) (Figure 1). Gangliosides are embedded in the outer layer of the plasma membrane and are one of major components of lipid rafts.⁷ Therefore, alteration in the ganglioside composition of lipid rafts can affect the localization of lipid raft-associated proteins.^{11–13} For example, knockout (KO) mice lacking GD2/GM2 synthase, in which GA2, GM2, GD2, GT2, and downstream gangliosides are depleted, show the dispersion of lipid raft-associated proteins, such as caveolin 1 or flotillin 1, from lipid raft fractions in flotation assay.¹³ Furthermore, in KO mice lacking both GD2/GM2 synthase and GD3 synthase, a GPI-anchored complement-regulatory protein CD55/decay-accelerating factor also disperses from lipid rafts, and unfavorable activation of complement system occurs.¹² Complement activation starts with the activation of C1q by antigen-antibody complex or spontaneous hydrolysis of C3 and causes a cascade of further cleavage and activation processes of complement proteins. Complement activation results in the accumulation of complement proteins, such as C1q or C3b, and the formation of the membrane attack complex. In the KO mice lacking both GD2/GM2 synthase and GD3 synthase, gene expression of complement proteins, such as C1qa, C3, or C4, was up-regulated, and deposition of C1q in the brain increased.^{12,13} Meanwhile, an activation of the complement system and accumulation of C1q, C3, or membrane attack complex have also been demonstrated in brains of human patients with prion diseases or prion-infected animals.^{14,15} The role of complement activation in prion disease

pathogenesis has been proposed to cause degeneration of prion-infected neurons or to facilitate microglial phagocytosis of prion-infected neurons.¹⁵

In this study, we performed a prion transmission study using various ganglioside synthase knockout mouse strains to investigate the role of gangliosides in the pathogenesis of prion diseases.

Materials and Methods

Ethics Statement

Animal experiments were performed in strict accordance with the Regulations for Animal Experiments and Related Activities at Hokkaido University and Fundamental Guidelines for Proper Conduct of Animal Experiment and Related Activities in Academic Research Institutions by Ministry of Education, Culture, Sports, Science and Technology in Japan (notice number 71). The protocol was approved by the Institutional Animal Care and Use Committees of Hokkaido University (14-0170).

Mice

Generation of KO mice lacking GD2/GM2 synthase (B4GALNT1), GD3 synthase (ST8SIA1), or GM3 synthase (ST3GAL5) has been reported previously.^{16,17} These mice were backcrossed to C57BL/6, and the resulting heterozygous animals were intercrossed to provide homozygous KO mice and wild-type (WT) littermate controls. Thus, all of the KO mice and WT littermate controls were on a C57BL/6 background.

Figure 1 Biosynthetic pathway of gangliosides and elimination of particular gangliosides in KO mouse strains. GD2/GM2, GM3, and GD3 KO mice lack gangliosides, indicated by red, blue, and pink lines, respectively.

Sources of Inocula

A mouse-adapted scrapie prion strain Chandler¹⁸ was kindly provided by Dr. Motohiro Horiuchi (Faculty of Veterinary Medicine, Hokkaido University, Sapporo, Japan). A mouse-adapted Gerstmann-Sträussler-Scheinker syndrome prion strain Fukuoka-1¹⁹ was maintained in C57BL/6 mice. The inocula were prepared from the brain of terminally ill mice.

Transmission Experiments

Intracerebral inoculation was performed as described.²⁰ Briefly, 10% brain homogenates were prepared in sterile phosphate-buffered saline using glass homogenizers, and 20 μ L of the homogenates was intracerebrally inoculated into mice ($n = 6$ or 7). The inoculated mice were sacrificed at a predefined clinical end point or at the time point showing intercurrent illness. One hemisphere of the brain was fixed in 10% buffered formalin for histopathological analysis, and the other hemisphere was immediately frozen for biochemical analysis.

Immunohistochemistry

Formalin-fixed mouse brain tissues were treated with 60% formic acid for 1 hour to inactivate the infectivity and embedded in paraffin. The embedded tissues were divided into sections (5 μ m thick). Tissue sections were pretreated by hydrolytic autoclaving before PrP immunohistochemistry.²¹ The anti-PrP antiserum PrP-N,²² rabbit anti-ionized calcium-binding adapter molecule 1 (Iba1) polyclonal antibody (FUJIFILM Wako Pure Chemical Corp., Osaka, Japan), and rabbit anti-glial fibrillary acidic protein (GFAP) polyclonal antibody (Dako, Glostrup, Denmark) were used as the primary antibodies. Goat-anti-rabbit Ig polyclonal antibody labeled with the peroxidase-conjugated dextran polymer, EnVision⁺ (Dako), was used as the secondary antibody. For routine histopathological analysis, the tissue sections were also stained with hematoxylin and eosin.

PrP^{Sc} Purification

PrP^{Sc} was purified from the mouse brain as described.²³ Briefly, brain tissues were homogenized in 2 mL of lysis buffer [100 mmol/L tris-HCl, pH 8.0, 10 mmol/L NaCl, 10 mmol/L MgCl₂, 2% Triton X-100, and 25 U/mL DNase I (Takara Bio, Shiga, Japan)] and digested with collagenase (1 mg/200 mg tissue) (FUJIFILM Wako Pure Chemical Corp.) overnight at room temperature. Collagenase digestion disrupts the connective tissue and improves the accessibility of detergents and/or proteinase K (PK) to PrP^{Sc}.²⁴ The digested homogenates were ultracentrifuged at 453,000 $\times g$ for 30 minutes at 4°C, and the pellets were resuspended and sonicated in 870 μ L of PK-digestion buffer [100 mmol/L tris-HCl, pH 8.0, and 5% Sarkosyl (Sigma-

Aldrich, St. Louis, MO)]. The resuspended samples were centrifuged at 10,000 $\times g$ for 3 minutes to remove the cell debris, and the supernatants (800 μ L) were digested with PK (4 μ g/200 mg tissue) (FUJIFILM Wako Pure Chemical Corp.) for 1 hour at 37°C. It has been reported that these conditions for PK digestion were sufficient for the complete digestion of normal PrP^C and that higher PK concentrations caused unfavorable degradation of PrP^{Sc}.²⁵ The PK-digested proteins were precipitated by adding 200 μ L of 99.5% ethanol and ultracentrifugation at 135,000 $\times g$ for 30 minutes at 4°C. The pellets were resuspended in 400 μ L/200 mg tissue of Laemmli's sample buffer (60 mmol/L tris-HCl, pH 6.8, 5% glycerol, 2% SDS, and 0.01% bromophenol

Figure 2 Kaplan-Meier survival curves. Intracerebral transmission of Chandler (A) or Fukuoka-1 (B) strain to each KO mouse strain or wild-type (WT) littermate controls, respectively.

blue) and boiled at 100°C for 5 minutes. To examine the expression level of PrP^C, C1qa, and β -actin, 10% homogenates were prepared from uninfected mouse brains using the lysis buffer. The homogenates were boiled with the addition of 0.25× volume of 5× Laemmli's sample buffer.

Flotation Assay

To isolate lipid rafts, a flotation assay was performed as described previously.²⁶ Briefly, 10% homogenates were prepared from uninfected mouse brains using citrate buffer (20 mmol/L sodium citrate and 137 mmol/L NaCl, pH 6.0). The homogenates were centrifuged at $1000 \times g$ for 5 minutes at 4°C to remove cell debris and nuclei. The supernatants were mixed with 1× volume of ice-cold Brij O-10 extraction buffer [0.1% Brij O-10 (Sigma-Aldrich), 20 mmol/L sodium citrate, and 137 mmol/L NaCl, pH 6.0] and incubated on a rocker at 4°C for 30 minutes, and then mixed with 1.4× volume of 60% Optiprep (Axis-Shield, Dundee, UK). In 12 mL polycarbonate centrifuge tubes (Beckman Coulter, Brea, CA), 9 mL of 0% to 30% Optiprep continuous gradients were prepared by Gradient Master (BioComp Instruments, Frederickton, NB, Canada), according to the manufacturer's instruction, and underlaid with 2 mL of the Brij O-10 extracted samples. The gradients were centrifuged in a prechilled Beckman SW41 rotor (Beckman Coulter) at $40,000 \times g$ for 130 minutes at 4°C. The samples were split to 18 fractions (600 μ L each) from top (fraction 1) to bottom (fraction 18) and boiled with the addition of 150 μ L of 5× Laemmli's sample buffer.

Western Blot Analysis

Protein samples were subjected to SDS-PAGE using 13.5% tris-glycine long gels of 15 cm length and Western blot analysis, as described.²⁷ The anti-PrP monoclonal antibody SAF83 (Bertin Bioreagents, Montigny le Bretonneux, France), anti- β -actin monoclonal antibody 2F3 (FUJIFILM Wako Pure Chemical Corp.), anti-flotillin 1 monoclonal antibody 18/Flotillin-1 (BD Biosciences, San Jose, CA), anti-C1qa polyclonal antibody (Abcam, Cambridge, UK), anti-CD55 monoclonal antibody number 583905 (R&D Systems, Minneapolis, MN), anti-voltage-gated calcium channel auxiliary subunit $\alpha 2\delta$ -1 (CACNA2D1) monoclonal antibody 20A (Abcam), and anti-Thy1.2 monoclonal

antibody (Bio-Rad, Hercules, CA) were used as the primary antibodies. The anti-mouse EnVision+ (Dako), anti-rabbit EnVision+ (Dako), and electrochemiluminescent anti-rat IgG antibody (GE Healthcare, Chicago, IL) were used as the secondary antibodies. The blots were visualized with Luminata Forte Western HRP substrate (MilliporeSigma, Burlington, MA), and images were obtained by imaging device ImageQuant LAS 4000 mini (GE Healthcare). The signal intensities of the Western blot analyses were quantified with ImageQuant TL software version 7.0 (GE Healthcare).

Image Analysis

Severity of spongiform change was classified into three scores: 1, negative/mild (approximately 0% to 20% of the tissue section showed spongiform change); 2, moderate (approximately 20% to 60% of the tissue section showed spongiform change); and 3, severe (approximately 60% to 100% of the tissue section showed spongiform change). Immunohistochemical staining intensities were classified into three scores: 1, negative/weak (approximately 0% to 20% of the tissue section showed immunoreactivity); 2, moderate (approximately 20% to 60% of the tissue section showed immunoreactivity); and 3, high (approximately 60% to 100% of the tissue section showed immunoreactivity). Cerebral cortex, cerebellar cortex, striatum, thalamus, hippocampus, pons, and cerebellar white matter sections were used for the scoring of spongiform change, and cerebral cortex sections were used for the scoring of immunohistochemistry. Scoring was performed independently by two pathologists in a blinded manner (A.K. and Z.Q.), and the scores were finalized after discussion. Data were collected from all diseased mice and are represented as means $\pm$ SEM in the scoring of spongiform change or as percentage of animals with each score in the scoring of immunohistochemistry.

Statistical Analysis

The Kaplan-Meier log-rank test was used to analyze survival data in transmission study. Tukey-Kramer test was used to analyze signal intensity data in Western blot analysis. The statistical tests were performed using the statistical software EZR version 1.36.²⁸ $P < 0.05$ was considered statistically significant.

Figure 3 Neuropathological features. **A:** Histopathological analysis of spongiform changes in hematoxylin and eosin (H&E)-stained sections and immunohistochemical analysis for PrP^{TSE} deposition, ionized calcium-binding adapter molecule 1 (Iba1)-positive microglia, and glial fibrillary acidic protein (GFAP)-positive astroglia in the cerebral cortices of the Chandler strain-inoculated mice. GD2/GM2 KO mice showed less PrP^{TSE} deposition in the neuropil and attenuated microglial and astroglial reactions. Data are representative of tissue sections from six mice per group. **B:** Histopathological analysis of the Fukuoka-1 strain-inoculated mice. **C:** Regional distribution of spongiform change in the brain. Spongiform change was scored on a scale of 1 to 3: 1, negative/mild (approximately 0% to 20% of the tissue section showed spongiform change); 2, moderate (approximately 20% to 60% of the tissue section showed spongiform change); and 3, severe (approximately 60% to 100% of the tissue section showed spongiform change). **D:** Scoring of the immunohistochemical signal intensities. The signal intensities were classified into three scores: 1, negative/weak (approximately 0% to 20% of the tissue section showed immunoreactivity); 2, moderate (approximately 20% to 60% of the tissue section showed immunoreactivity); and 3, high (approximately 60% to 100% of the tissue section showed immunoreactivity). Data are expressed as percentage of animals with each score. The scoring was performed by two pathologists independently (A.K. and Z.Q.), on the basis of the signal intensities in the cerebral cortices. Data are expressed as means $\pm$ SEM (**C**). $n = 6$ or 7 (**C** and **D**). Scale bars = 50 μ m (**A**). WT, wild type.

Figure 4 Western blot analysis of PrP. **A:** Proteinase K (PK)–resistant PrP^{Sc} in the brain of Chandler- or Fukuoka-1–inoculated mice. There was no difference in the biochemical properties of PK-resistant PrP^{Sc} among mouse strains. The Chandler-inoculated mice consistently produce lower apparent molecular mass of unglycosylated PrP^{Sc} fragments compared with those of the Fukuoka-1–inoculated mice regardless of their genotype. The Chandler-inoculated GD2/GM2 KO mice showed slightly decreased apparent amount of PrP^{Sc} compared with wild-type (WT), GD3 KO, or GM3 KO mice, irrespective of the degree of PrP^{TSE} deposition in the immunohistochemical analysis. For example, a GD2/GM2 KO mouse (number 145) that showed reduced PrP^{TSE} deposition in the immunohistochemical analysis (ie, score 1) has similar apparent amount of PK-resistant PrP^{Sc} compared with that of another GD2/GM2 KO mouse (number 203) that shows abundant PrP^{TSE} deposition in the immunohistochemistry (ie, score 3). A brain sample, equivalent to 125 μ g in wet weight, was loaded in each lane. **B:** Quantification of the amount of PK-resistant PrP^{Sc} in the brain of the Chandler-inoculated mice. GD2/GM2 KO mice show reduced PrP^{Sc} accumulation compared with GD3 KO mice or GM3 KO mice. The mean signal intensity of PrP^{Sc} in GD2/GM2 KO mice was assigned as 100. Statistical significance calculated by Tukey–Kramer test. **C:** Expression levels of PrP^C in the brain of uninfected mice. A brain sample, equivalent to 50 μ g in wet weight, was loaded in each lane. Data are expressed as means $\pm$ SEM (**B**). $n = 4$ or 5 (**B**). * $P < 0.05$.

Results

Transmission of Prions to KO Mice Lacking Ganglioside Synthase Gene

To investigate the role of gangliosides in the pathogenesis of prion diseases, intracerebral transmission study was performed using KO mice lacking GD2/GM2 synthase (GD2/GM2 KO), GD3 synthase (GD3 KO), or GM3 synthase (GM3 KO). These KO mice lack particular ganglioside species, depending on the ablated ganglioside synthase gene (Figure 1). For a mouse-adapted scrapie strain Chandler, the mean incubation time of GD2/GM2 KO mice (126.6 ± 0.3 days) was reduced by 20% compared with that of WT mice (157.5 ± 1.2 days; $P < 0.01$) (Figure 2). The mean incubation time of GM3 KO mice was also slightly reduced (149.5 ± 2.2 days; $P < 0.01$). For a mouse-adapted Gerstmann–Sträussler–Scheinker syndrome strain Fukuoka-1, the mean incubation time of GD2/GM2 KO mice and GM3 KO mice was reduced compared with that of WT mice, although the differences were not statistically significant.

Neuropathology

Prion strains are characterized by neuropathological features and biochemical properties of PrP^{Sc} in the inoculated mice. Therefore, neuropathological analysis of the inoculated mice was performed to examine whether the strain characteristics were altered by the ablation of the ganglioside synthase gene. In the Chandler-inoculated mice,

spongiform changes were distributed through the cerebral and cerebellar cortices, striatum, thalamus, hippocampus, pons, and cerebellar white matter (Figure 3A). In the Fukuoka-1–inoculated mice, the vacuoles of spongiform changes became larger and were distributed more intensively in the pons and cerebellar white matter (Figure 3, B and C). There was no difference in the distribution of spongiform changes among mouse strains (Figure 3C). However, in immunohistochemical analysis of disease-associated PrP, PrP^{TSE}, GD2/GM2 KO mice inoculated with the Chandler strain showed decreased PrP^{TSE} deposition (Figure 3, A and D). In addition, Iba 1–positive microglia and GFAP-positive astroglia were also fewer in the Chandler-inoculated GD2/GM2 KO mice. For the Fukuoka-1 strain, GD2/GM2 KO mice showed decreased

Figure 5 Activation of complement systems. Western blot analysis of C1qa in the brain of uninfected mice and Chandler- or Fukuoka-1–inoculated mice. C1qa accumulated only in the prion-infected animals regardless of the mouse genotype or prion strain. A brain sample, equivalent to 1 mg in wet weight, was loaded in each lane. WT, wild type.

Figure 6 Flotation assay of lipid raft-associated proteins. Floating patterns of a glycosylphosphatidylinositol (GPI)-anchored complement-regulatory molecule CD55 (**A**), a lipid raft marker flotillin 1 (**B**), PrP (**C**), a GPI-anchored PrP-associated calcium channel protein CACNA2D1 (**D**), and a GPI-anchored protein Thy1 (**E**) in fractions separated by Optiprep density gradient centrifugation were examined using brain homogenates of uninfected mice. Flotillin 1 and PrP dispersed from the lipid raft fractions (fractions 7 to 11) to the nonraft fractions (fractions 15 to 18) in GD2/GM2 KO mice. Data are representative of assays repeated independently three times. WT, wild type.

microglial reaction, but PrP^{TSE} deposition or astroglial reaction was not altered (**Figure 3, B and D**).

Biochemical Properties of PrP^{Sc}

Western blot analysis of PK-resistant PrP^{Sc} in the brain revealed that the strain-specific biochemical properties of

PrP^{Sc} (ie, apparent molecular mass and patterns of *N*-linked glycosylation) were not modified by the absence of ganglioside synthase genes (**Figure 4A**). The apparent molecular mass of PrP^{Sc} in the Chandler-inoculated mice was smaller than that in the Fukuoka-1-inoculated mice, regardless of the mouse genotype. Meanwhile, the Chandler-inoculated GD2/GM2 KO mice showed slightly

decreased PrP^{Sc} accumulation compared with WT, GD3 KO, or GM3 KO mice (Figure 4, A and B). The apparent amount of PrP^{Sc} in Western blot analysis did not reflect the degree of PrP^{TSE} deposition in the immunohistochemical analysis. For example, a GD2/GM2 KO mouse (number 145) that showed reduced PrP^{TSE} deposition in the immunohistochemical analysis (ie, score 1) had similar apparent amount of PrP^{Sc} to that of another GD2/GM2 KO mouse (number 203) that showed abundant PrP^{TSE} deposition in the immunohistochemistry (ie, score 3) (Figure 4A). Thus, GD2/GM2 KO showed shortened incubation time, reduced PrP^{TSE} deposition with attenuated glial reactions in neuropathological analysis, and slightly decreased PK-resistant PrP^{Sc} accumulation in Western blot analysis.

In addition, the amount of PrP^C was compared in the brain of uninfected KO mice because the expression level of PrP^C can directly affect the incubation period. However, the amount of PrP^C was not modified by the absence of ganglioside synthase genes (Figure 4C), suggesting that the expression level of PrP^C is unrelated to the changes in incubation period.

Evaluation of Complement System

To gain insights into the shortened incubation period in GD2/GM2 KO mice, the involvement of complement system was examined because it has been reported that KO mice lacking both GD2/GM2 synthase and GD3 synthase show dispersion of GPI-anchored complement-regulatory proteins from lipid rafts and activation of the complement system.^{12,13} To assess the activation of the complement system, the amount of a complement factor C1qa in the brain was compared between GD2/GM2 KO mice and WT mice by Western blot analysis (Figure 5). However, there was no difference in C1qa level between GD2/GM2 KO mice and WT mice. C1qa accumulated only in the prion-infected animals regardless of the mouse genotype or prion strain. In addition, although dispersion of GPI-anchored complement-regulatory protein CD55 from lipid rafts has been demonstrated in KO mice lacking both GD2/GM2 synthase and GD3 synthase, the floating patterns of CD55 were not different between GD2/GM2 KO mice and WT mice in the flotation assay of lipid raft-associated proteins in the brain (Figure 6A), suggesting that CD55 did not disperse from lipid rafts in GD2/GM2 KO mice. Thus, the shortened incubation period in the Chandler-inoculated GD2/GM2 KO mice was not due to altered activation of the complement system.

Flotation Assay of PrP^C and Lipid Raft–Associated Proteins

It was subsequently investigated whether PrP^C and lipid raft-associated proteins other than CD55 disperse from lipid rafts in GD2/GM2 KO mice by flotation assay. As reported previously,¹³ a lipid raft marker, flotillin 1,

dispersed from the lipid raft fractions (fractions 7 to 11) to the nonraft fractions (fractions 15 to 18) in GD2/GM2 KO mice (Figure 6B). Furthermore, PrP^C also dispersed from the lipid raft fractions to the nonraft fractions in GD2/GM2 KO mice (Figure 6C). In contrast, the floating patterns of a GPI-anchored PrP-associated calcium channel protein CACNA2D1^{29,30} or those of a GPI-anchored protein Thy1 were not different between GD2/GM2 KO mice and WT mice (Figure 6, D and E). Thus, certain lipid raft-associated proteins, including PrP^C, dispersed from lipid rafts in the brain of GD2/GM2 KO mice.

Discussion

Herein, we found that GD2/GM2 KO mice inoculated with the Chandler prion strain showed shortened incubation time, reduced PrP^{TSE} deposition with attenuated glial reactions in neuropathological analysis, slightly decreased PK-resistant PrP^{Sc} accumulation in Western blot analysis, and reduced localization of PrP to lipid rafts.

The reduced localization of PrP to lipid rafts might affect the formation of PrP^{Sc} in GD2/GM2 KO mice. Although the localization of PrP^{Sc} to lipid rafts could not be examined, dispersion of lipid raft-associated proteins, including PrP^C from lipid rafts, was confirmed in GD2/GM2 KO mice in the present study. Because the localization to lipid rafts is important for the conformational conversion of PrP^C into PrP^{Sc},³ the dispersion of PrP^C and/or PrP^{Sc} from lipid rafts might result in the decreased accumulation of PrP^{TSE} and PK-resistant PrP^{Sc} in the Chandler-inoculated GD2/GM2 KO mice. The amount of PK-resistant PrP^{Sc} in the Chandler-inoculated GD2/GM2 KO mice in the Western blot analysis did not reflect the marked reduction of PrP^{TSE} deposition in the immunohistochemical analysis compared with the other mouse strains. In addition, even in the Chandler-inoculated GD2/GM2 KO mouse group, the apparent amount of PK-resistant PrP^{Sc} did not reflect the PrP^{TSE} deposition score in the immunohistochemical analysis. This finding raises a possibility that the amount of PK-sensitive PrP^{Sc}^{31–33} might largely decrease in the Chandler-inoculated GD2/GM2 KO mice. Meanwhile, the incubation time was shortened in the Chandler-inoculated GD2/GM2 KO mice despite the decreased PrP^{Sc} accumulation. To account for this discrepancy, further studies will be needed in the future addressing the questions of whether the dispersion of PrP^{Sc} and/or PrP^C from lipid rafts reduces the formation of PK-sensitive PrP^{Sc} or affects the neurotoxicity of PrP^{Sc}.

There remains a possibility that the alteration in ganglioside composition might directly affect the conformation of PrP. The ablation of ganglioside synthase causes massive alteration of ganglioside composition.¹⁷ For example, an ablation of GD2/GM2 synthase causes depletion of GD2, GM2, and downstream gangliosides, such as GM1, and results in accumulation of GD3 and GM3.^{16,34} PrP^C is

colocalized and coimmunoprecipitated with GM1 and GM3.^{35,36} GM1 binds to the C-terminus of PrP and modifies the conformational properties of PrP.^{37,38} Similarly, GM1 binds to amyloid β and modifies the conformation of amyloid β , resulting in the formation of endogenous seeds for amyloid fibril formation.³⁹ Therefore, the direct effects of gangliosides on PrP conformation are not negligible. To test this hypothesis, analysis of conformational properties of PrP^C and PrP^{Sc} from GD2/GM2 KO mice will be needed in the future.

A possibility that the alteration in ganglioside composition might affect the activation of glial cells cannot be excluded. Although the role of glial cells in the pathogenesis of prion diseases is controversial, both microglia and astroglia are activated in prion diseases.^{40–43} The decreased number of microglia and astroglia in the Chandler-inoculated GD2/GM2 KO mice might simply reflect the decreased PrP^{TSE} deposition. However, although the amount of PrP^{TSE} deposition was not altered, the Fukuoka-1-inoculated GD2/GM2 KO mice also showed attenuated microglial reaction. Therefore, the altered ganglioside composition may affect glial reactions regardless of the PrP^{TSE} accumulation. Indeed, ethanol-induced astroglial activation is attenuated in GD2/GM2 KO mice.⁴⁴ Further studies will be required to examine whether the shortened incubation time in GD2/GM2 KO mice relates to the attenuated microglial reaction, including neuroprotective microglia.^{45,46}

In conclusion, we have shown that the ablation of GD2/GM2 synthase gene causes a significant reduction in incubation time of prion infection. The present study suggests that the gangliosides may have an important role in prion disease pathogenesis by affecting the localization of PrP to lipid rafts.

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Supplemental Data

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