**Exploration of the main sites for the transformation of normal prion protein (PrP<sub>C</sub>) into pathogenic prion protein (PrP<sub>Sc</sub>)**

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**Abstract**

**Introduction:** The functions and mechanisms of prion proteins (PrP<sub>C</sub>) are currently unknown, but most experts believe that deformed or pathogenic prion proteins (PrP<sub>Sc</sub>) originate from PrP<sub>C</sub>, and that there may be plural main sites for the conversion of normal PrP<sub>C</sub> into PrP<sub>Sc</sub>. In order to better understand the mechanism of PrP<sub>C</sub> transformation to PrP<sub>Sc</sub>, the most important step is to determine the replacement or substitution site. **Material and Methods:** BALB/c mice were challenged with prion RML strain and from 90 days post-challenge (dpc) mice were sacrificed weekly until all of them had been at 160 dpc. The ultra-structure and pathological changes of the brain of experimental mice were observed and recorded by transmission electron microscopy. **Results:** There were a large number of pathogen-like particles aggregated in the myelin sheath of the brain nerves, followed by delamination, hyperplasia, swelling, disintegration, phagocytic vacuolation, and other pathological lesions in the myelin sheath. The aggregated particles did not overflow from the myelin in unstained samples. The phenomenon of particle aggregation persisted all through the disease course, and was the earliest observed pathological change. **Conclusion:** It was deduced that the myelin sheath and lipid rafts in brain nerves, including axons and dendrites, were the main sites for the conversion of PrP<sub>C</sub> to PrP<sub>Sc</sub>, and the PrP<sub>Sc</sub> should be formed directly by the conversion of protein conformation without the involvement of nucleic acids.

**Keywords:** mice, brain, prion proteins, myelin sheath, particle aggregation.

**Introduction**

Prion diseases predominantly affect the nervous system leading to fatal and untreatable degenerative disorders of the brain, which are called transmissible spongiform encephalopathies (TSEs). The causative agent of TSEs has been identified as a prion known to be an abnormal isoform (Scrapie prion protein, PrP<sub>Sc</sub>) of a normal cellular prion protein (PrP<sub>C</sub>) (5). PrP<sub>C</sub> is found in all cells of the body, with relatively high levels in the brain. PrP<sub>Sc</sub> is able to induce transformation of the normal prion protein nearby into the misfolded and infectious prion (23).

PrP<sub>Sc</sub> differs from PrP<sub>C</sub> neither in its covalent structure nor post-translational modification, but only in its conformation. PrP<sub>Sc</sub> mainly has α-sheeted (helical or β-sheet) conformation, while PrP<sub>C</sub> chiefly shows helical conformation (6). A substantial proportion of investigators believe that PrP<sub>Sc</sub> (an abnormal conformer) can convert PrP<sub>C</sub> (the normal cellular protein) into PrP<sub>Sc</sub>. This transformation process may be accomplished through PrP<sub>Sc</sub> serving either as a template or a seed to induce transformation of PrP<sub>C</sub> (4). Hence, prion diseases belong to a wide group of conformational disorders or diseases of protein misfolding. The prion theory is almost universally accepted nowadays and its author has been honoured by the Nobel prize (31), although the theory remains to be formally proved. Great success has been achieved in research, yet it is still difficult to explain the reason for variation in scrapie (10).
Furthermore, there are various clarification theories pertaining to the mechanism behind how and where PrPSc induces the transformation of PrPC. Some pathological phenomena caused by prion diseases like vacuolisation and myelin sheath puff region changes may at least in part originate from PrPSc accumulation (34), and several studies have speculated that myelin shear cytoplasmic membrane/sterol rafts of nerve fibres and endoplasmic reticulum may be the main conversion sites of PrPSc into PrPSc. Irrespective of whether there is or is not an association with the PrPSc accumulations, it is still important to clarify early pathogenetic phenomena. The mechanism of the conversion conversion of prion also involves relative special environments, various relative elements, and predisposition factors (25). Nonetheless, how the exogenous PrPSc enters the brain and central nervous system along the nerve after the infection of the peripheral nervous system with prion is still a mystery. In general, it is believed that the prion protein passes through a nerve fibre. However, the particular transformation mechanism is not clear (3, 18, 28). PrPC is attached to the exterior of the plasma membrane by its glycosyl-phosphatidyl-inositol (GPI) anchor. Currently, a large number of ligands are known to interact with PrPC that have also been implicated in a wide range of cellular functions. The way in which PrPC is anchored to the membrane plays an important role in turnover and pathogenesis of PrPSc. GPI anchors do not extend beyond the top leaflet of the lipid bilayer. Therefore, the proteins such as PrPC must be associated with proteins containing transmembrane sections for them to be able to communicate in the intercellular processes. Current data suggest that the clustering of PrPC into distinct membrane patches may be due to the association of PrPC with transmembrane proteins responsible for trafficking PrPSc into endocytic pathways (15). Moreover, involvement of transmembrane proteins in the pathological conversion of PrPC into PrPSc remains unknown.

In this study, we observed the ultra-structure and pathological changes of the brain of mice infected with prion strain RML to define the mechanism of conformation changes of prion protein, the substitution sites, and the conditions for the transformation of the prion protein into the pathogenic form. The lesions in myelin shear of nerve fibres including dendrites and axons were investigated to know how PrPC converts into PrPSc at the main substitution sites of PrPC.

Material and Methods

Prion strain source and experimental animals. The prion strain RML was kindly provided by the China Animal Health & Epidemiology Center, National Foreign Animal Disease Diagnostic Center (Qingdao). Female BALB/c mice (about 21 g b.w.) were raised in cages with occupancy of five and abundant supply of feed and water at a temperature of 22°C ±2°C and relative humidity of 55% ±5% under natural daylight conditions.

Groups and treatment of animals. There were 70 mice divided into four groups. The test group consisted of 55 mice challenged with mouse-adapted prion strain RML. The three control groups included a blank control (five mice) treated with physiological saline (0.85% NaCl), an experimental control (five mice) with brain homogenates of normal mice, and a normal control (five mice) without any treatment. The brain homogenates of mice infected with prion strain RML were prepared as mouse-adapted prion strain RML, which was diluted 10-fold with physiological saline as prion challenge solution. After anaesthesia, 25 μL of prion challenge solution was injected into the brains of mice in the test group. Meanwhile, mice in the blank control group and experimental control group were respectively injected with the same amount of physiological saline or brain homogenates of normal mice by the same injection method. Changes in the behaviour and body weight of the mice were regularly observed and recorded.

Sampling of mouse brain. From 90 dpc (when the primal nervous symptoms start to appear), the brains of five mice in the test group were sampled once a week and immediately fixed in 2.5% glutaraldehyde buffer (0.1 M sodium phosphate, pH 7.4) for ultra-structure observation. Brain samples from the five mice in each of the control groups were collected at 160 dpc, and handled similarly to the test group.

Ultra-structure observation of mouse brain under transmission electron microscope (TEM). Preparation of pathological slices from samples of fixed mouse brain was carried out according to previous reports (13, 16). In brief, about 2 mm³ of brain samples for TEM were immersed in 2.5% glutaraldehyde buffer (pH 7.4 with 0.1 M sodium phosphate) and kept at 4°C for more than 24 h, and then washed three times in cold sodium phosphate buffer. Later, the brain tissues were post-fixed in 1% osmium tetroxide solution for 2 h at 4°C. Following that, the fixed tissues were dehydrated on an increasing step ethanol gradient, sunk into propylene oxide for 60 min, and embedded in SPI-PON812 kit (SPI Supplies, USA) for 12 h according to the instructions of manufacturer. Ultrathin sections were cut using a Leica UC7 ultramicrotome (Leica, Germany) and stained with uranyl acetate and lead citrate. Finally, the ultra-structure of the brains was observed with a Hitachi H-7650 TEM (Hitachi, Japan) at 80 kV.

Results

Observation of mental status and behaviour. The mice started to present clinical symptoms from 90 days after RML challenge, and the main symptoms were depression, autism, loss of hair lustre, and ataxia. With the progress of the infection, lethargy and sensory disturbance appeared in the late period.
Observation of myelinated nerves in the control groups. As shown in Fig. 1A, the structures of myelinated nerves from the normal control group at 160 dpc were compact, the myelin membrane and myelin were closely connected without separation, and there was no stratification in the myelin. Myelinated vacuoles could be seen occasionally and the subcellular structures like mitochondria were integrated in the sheath. Structural abnormality of myelinated nerves appeared neither in the blank control group (Fig. 1B) nor in the experimental control group (Fig. 1C) at 160 dpc just as it did not in the normal control.

Observation of myelinated nerves in the test group from 90 to 160 dpc. Changes of myelin were shown at 90 dpc. The vacuolar separation had become larger in myelin and increased gradually, more and more caveolae had emerged on the surface of myelin sheaths with tension reduction, and stratification and separation between the cytoplasmic membrane and myelin had also begun to take place. In addition, deposition of dispersed black particles (uncertain pathogens) had increased, and become very small. The myelin had begun to deform and the structure of myelin sheaths was fuzzy. These changes were obvious especially between the two nerve fibres (Fig. 2A). As seen in Fig. 2B, it was also found that local structure of myelin had become blurred and disintegrated, and the original inherent structure of myelin had almost disappeared. Some of the black depositions had accumulated into larger particles. Furthermore, the detached myelin plasmalemma was shrunken and detached vacuoles were also observed. However, the intracellular structure including organelles had not changed obviously, and only parts of the cytoplasmic membrane and myelin sheath were detached.

Fig. 1. The myelinated nerves of mouse brain from three control groups including the normal control (A), the blank control (B), and the experimental control (C). The normal control received no treatment, the blank control was treated with physiological saline (0.85% NaCl), and the experimental control was treated with brain homogenates of normal mice. The myelin sheaths in these control groups were compact without stratification in the myelin, and subcellular structures like mitochondria were integrated.

Fig. 2. Myelinated nerves of mouse brain from the test group at 90 days post challenge. Rightward arrows indicate structures of myelin sheath between two nerve fibres which had become fuzzy and deformed, a leftward arrow indicates that there were some black depositions accumulated into larger particles, and a downward arrow indicates the detached invaginated myelin plasmalemma and vacuoles.
At 97 dpc, the myelin had become curved with tension markedly changed to looseness and there was apparent stratification in the myelin, clear deposition or polymerisation of dispersed particles, and an unintegrated cytoplasmic membrane (Fig. 3A). The structure of mitochondria was almost intact, but some parts of the mitochondrial membrane had broken down, and were tending toward degeneration. The stratification in the myelin sheath was more obvious where there were black particles accumulated (Fig. 3B). The myelinated fibres were dissociated, and the inherent structure was destroyed, by which to say it had a swollen state as compared to the structure of local myelin. Moreover, black particles were only aggregated in the myelin.

Some pathogenic changes in the brain at 104 dpc are shown in Fig. 4. Deposition or polymerisation of the particles was much more evident, yet they were still limited within the myelin sheath. Some part of the myelin had dissolved and the damage was more significant, including lysis of the plasma membrane (Fig. 4A, rightward arrow). Bubbly structures (Fig. 4A, leftward arrow) had appeared, yet organelle damage was not obvious. Degeneration was more likely to have happened at the border between two myelin sheaths (Fig. 4B, rightward arrow), and adjacent mitochondria were also damaged due to deterioration of the myelin (Fig. 4B, downward arrow).

**Fig. 3.** The myelinated nerves of mouse brain from the test group at 97 days post challenge. The myelin had become curved and deposition or polymerisation of dispersed particles was obvious in the myelin sheath. Rightward arrows indicate that the cytoplasmic membrane was no longer integrated and leftward arrows indicate that stratification in the myelin sheath was more obvious where there were black pathogen-like particles accumulated.

**Fig. 4.** The myelinated nerves of mouse brain from the test group at 104 days post challenge. The plasma membrane had dissolved and spread out like mud (rightward arrow). In addition, there were bubbly structures (leftward arrow). Degenerative changes at the border between two myelin sheaths were more obvious (rightward arrow) and adjacent mitochondria are also damaged (downward arrow).
At 111 dpc, the cytoplasmic membrane was separated or destroyed, myelin detached from the main body, and there were large black particles accumulated. Furthermore, the polymeric particles were also present in the polymerised bubbles along with the detached myelin sheath (Fig. 5A, downward arrow). The myelin was disintegrated and scattered in the medullary cavity where some black particles were also aggregated. The structure of mitochondria was almost normal, bubbly structures had appeared once again (Fig. 5B), and there were suspicious pathogenic fibres (Fig. 5A, upward arrow).

At 118 dpc, pathological changes had occurred mainly in the myelin sheath (Fig. 6). It had become curved and a multilayer fiber layer had formed. Stratification was one of the significant characteristics in the local myelin sheath along with the aggregation of black particles. These deposited particles were bound with the solitary fibrous layer, and no free small and uniform black particles were observed. The structure of the myelin sheath was still integrated, with a slightly fuzzy appearance, and mitochondria were roughly intact. However, the myelin sheath had separated from the plasma membrane and a large gap had formed between two myelin sheaths. The aggregated black particles were mainly located in the lamination of the myelin sheath, and whether these black particles could become obvious seemed to be associated with the stratification extent. From 118 dpc, this stratification effect gradually diffused to both sides of the myelin sheath. The cytoplasm was like silk-slurry, while some specimens showed multiple sets of annular structure. Conformation changes in the myelin did not appear simultaneously, but usually these changes were more obvious in the adjacent dendritic myelin. There were widespread abnormal pathogenic structures at 118 dpc such as local extension, bending, and structural damage in the myelin sheath (Fig. 6). There were also many membrane cavities (autophagy vesicles) containing a membrane vacuole (Fig. 6A). Small black or specific particles were scattered among cells and tubulovesicular structures could also be seen (Fig. 6B).

**Fig. 5.** The myelinated nerves of mouse brain from the test group at 111 days post challenge. The rightward arrow points to the bubbly structures, the upward arrow points suspicious pathogenic fibres, and the downward arrows point to the polymerised bubble with polymeric particles.

**Fig. 6.** The myelinated nerves of mouse brain from the test group at 118 days post challenge. Some abnormal pathogenic structures like local extension, bending, and black particle deposition were widespread in the myelin sheath. The downward arrow indicates membrane vacuole, while the rightward arrow indicates tubulovesicular structure.
At 125 dpc, stratification in the myelin sheath had increased and become a universal abnormal structure. In addition, some myelin sheaths had become swollen and diffused (Fig. 7) and more particles had been deposited in local myelin. However, pathological changes were still limited to the myelin sheath and had rarely spread to the cytoplasm.

At 132 dpc, particle aggregation and stratified myelin were commonly observed features. The aggregated particles were distributed uniformly in the myelin sheath. The collapsed myelin sheath had spread to the cytoplasm and the infolded myelin sheath had further developed into phagocytic vacuoles (Fig. 8A). The typical Wallerian degeneration and intramyelinic vacuoles were obvious in the myelin sheath, while some myelin sheaths had formed multi-membrane vacuoles (Fig. 8B), in which the aggregated black particles were gradually dissolving. In the normal myelin, the aggregated black particles were not found, and the structures of cytoplasm and organelles were barely changed.

At 139 dpc, most of the dendrite myelin sheaths had swollen and become stratified, and particle aggregation or deposition was much more obvious than before (Fig. 9). The aggregated dark particles were gradually dissolving and further developing into a multivacuole structure, in which nothing tangible could be seen. Vacuoles or large areas of dissolved plaques had appeared outside the myelin. Polymeric particles attached to the myelin fibres were still visible. It was found that one myelin sheath contained another (Fig. 9A), and mitochondria had swollen conspicuously. In addition, both particle polymerisation and myelin sheath lamination had sharply increased (Fig. 9B). Although myelin still maintained its original shape, these changes had affected the basic form of cytoplasm, which had partly or completely dissolved. The black particles in the myelin sheath had diffused into the cytoplasm in relatively free forms. External tissues of the myelin sheath also showed the dissolved plaques.

At 146 dpc, the dark particle aggregations were more obvious in the myelin sheath, and some polymerised particles were in free forms, while most of them were attached to the myelin fibres. The cytoplasm was fully dissolved and multilayer membrane vesicles (large autophagy vacuoles) could also be seen. The whole myelin sheath cavity was filled with vacuoles and there was abundant proliferation of layered myelin fibres. The polymeric particles had become larger and there was myelin in a sleeve type or even in a four-layer casing pipe (Fig. 10, downward arrow). Black polymeric particles had increased obviously and were deposited in the myelin. The intracellular mitochondrial structure was integrated, but the myelin sheath was separated from the membrane (Fig. 10, upward arrows). It can be conclusively asseverated that the pathological changes in the brain infected with PrPSc first occur in the myelin sheath, and then in cytoplasm or external tissues.
Fig. 9. The myelinated nerves of mouse brain from the test group at 139 days post challenge. The myelin sheath is stratified obviously and many more particle aggregations or depositions were observed. The upward arrow indicates that one myelin sheath contained another downward arrows indicate particle polymerisation, and rightward arrows indicate myelin sheath lamination.

Fig. 10. The myelinated nerves of mouse brains from the test group at 146 days post challenge. The whole myelin sheath cavity is filled with vacuoles and layered myelin fibres, and there is typical particle deposition, myelin stratification, and vacuolisation. The downward arrow indicates the myelin in the form of a four-layer pipe, upward arrows indicate the myelin sheath separated from the membrane, rightward arrows indicate particle polymerisation and deposition, and leftward arrows indicate myelin sheath lamination. 4000×

At 153 dpc, the myelin sheath was further stratified and black particles much more aggregated. The cytoplasm had completely dissolved and separated from the membrane, and thus gave the appearance of vacuolisation. The axonal membrane had broken away from the myelin or proliferated to form an inner tongue (Fig. 11).

At 160 dpc, myelin stratification was still in progress and myelin had become curved due to the lack of tension (Fig. 12). The myelin detachment had caused vacuolisation of cytoplasm while the disaggregation of local myelin had led to the production of multiple vesicles. Almost all the myelin sheath was stratified and had swollen and become multi-hierarchical. Moreover, lesions were diffused outward, and part of the cytoplasm and extra-cellular structures had dissolved to form plaques. The mitochondria had swollen and become fuzzy, and adjacent structures of the cytoplasm and myelin were almost illegible. There were analogous autophagocytic vacuoles and multimembrane vesicles had appeared in the myelin.
sheath, large dissolved plaques were present (Fig. 12A), and the myelin close to plaques had disintegrated severely. With the continuous development of the pathological changes, polymeric particles had become more and more blurry, while the aggregated black particles in the myelin had almost dissolved to form plaque. Damage to the myelin had gradually expanded resulting in the disintegration and dissolution of myelin and cytoplasm respectively. However, most of the mitochondria still retained an intact structure. Membrane invagination and dissolved plaques exterior to the cytoplasm could also be seen. Nuclei of macrophagocytes were marginally assembled inside the cells (Fig. 12B). When the mice were on the verge of death, the nerve fibres had changed abnormally and inconsistently. However, stratification in the myelin sheath and aggregation of black particles were still existent. These black particles were considered universal pathological changes.

Discussion

Many potential roles and ligands for PrP<sup>C</sup> have been reported. Those studies suggest that PrP<sup>C</sup> may act as a scaffolding protein taking part in some cell surface interactions and signalling mechanisms. PrP<sup>C</sup>, an anchor protein (GPI) or a surface glycoprotein, is located on the extracellular surface and circulates between the plasma membrane and endocytic compartment. In addition, it participates in many biological processes of functional proteins, such as that of clathrin-plasma membrane associated caveolae-vesicle during the circulation process of cell membrane. Meanwhile, PrP<sup>C</sup> is an essential structure for the endocytosis of higher central nerve fibre content on the surface than other cells. The GPI of PrP<sup>C</sup> directly reacts with clathrin leading to cell invagination into the protoplasm. Several studies have implied that the conversion of PrP<sup>C</sup> to PrP<sup>Sc</sup> may take place during this process of invagination, and glycosylation of PrP<sup>C</sup> is the key to this misfolded transformation (29). In this transformation process, the exogenous PrP<sup>Sc</sup> is required as an incentive, and normal PrP<sup>C</sup> participates as the substrate to be transformed into an isoform or as a pathogenic factor (25). When this transforming process occurs in neuronal cells, it occurs mainly in the myelin sheath. Hence, the myelin sheath is the main site for prion conversion, which has been verified in the present study.

The normal structure of the myelin sheath of nerve fibres is compact with all laminations closely combined or the lipid rafts showing a laminated structure. The whole myelin is closely tied to the cytoplasmic membrane and the entire nerve fibre is orderly and tightly arranged. However, per observations of the brain pathological lesions of BALB/c mice infected with prion RML strain for 160 days, black particles first appeared in the myelin sheath of axons and dendrites or in lipid rafts. These black particles spread inside or outside the cell, and organelles such as the mitochondria were involved in further lesions that occurred in the myelin sheath. These findings indicated that the myelin sheath and lipid rafts were the first or main sites where the prion protein changed conformation (28). PrP<sup>C</sup> was expressed at a higher level in nerve cells especially in the myelin sheath, which may result from induction and serious pathological lesions caused by PrP<sup>Sc</sup> invasion. Besides this, PrP<sup>C</sup> was the main source of the substrate for abnormal transformation of PrP<sup>C</sup> induced by PrP<sup>Sc</sup>. Therefore, the myelin sheath with its high content of PrP<sup>C</sup> may become the main place where pathogenic PrP<sup>Sc</sup> is formed. In this study, we observed the lesions in mouse brain that were directly injected with prion proteins.
contained in brain homogenates. Results revealed that characteristic black particles were polymerised in the myelin sheath, which was not reported in previous studies. During the long period, all pathological lesions of the brain induced by PrPSc were continuously and systematically observed (1, 3). Different pathogenic strains would lead to different responses of the host (18). In the present study, prion strain RML was found to induce black particle aggregations in the myelin sheath, and in parallel with the advance of the infection, the deposition and aggregation of the black particles were much more obvious at successive intervals.

Some collapsed axons were surrounded by abnormal condensed multilamellar myelin sheaths in some nerve fibres, which strongly corresponded to an earlier study (3). Myelin sheaths with uniform or broken stratification were observed in the brain of mice injected with normal mouse brain homogenates and saline respectively, yet there were no black particles. The major structure of the myelin sheath is a lamination, while prion protein also has a lamellar conformation. PrPc is a membrane-bound glycoprotein, which mainly consists of an α-sheet layer and a β-sheet layer. However, the β-sheet of the lamellar structure is disordered as prion protein polymerisation occurs under pathological conditions (27).

It was supposed that the myelin sheath and plasma membrane were the key locations of PrPSc transformation as reported in earlier studies (18). According to the PrP endocytosis theory, PrPSc molecules first anchor on the surface of the membrane outside the myelin sheath, and then they may bend or invaginate when they encounter and bind with the myelin. For example, the myelin sheath becomes curved and its tensile strength is insufficient to keep it rigid and straight, as seen in the present and other studies (11). The sites undergoing bending or invagination were often the starting points of particle aggregations that subsequently led the myelin sheath to delaminate, as confirmed in many previous studies (26). Particle aggregations were the primary pathological lesions presented within the myelin sheath regardless of the structural changes in myelin. The black particles do not spread to the cytoplasm or outside the cell, possibly because the overflowing polymeric particles are dispersed or dissolved when they are in a different environment. Perhaps the potential locations for the conversion of PrPc to PrPSc are related to the myelin membrane, amyloid plaques, pathways from endocytosis to lysosomes, interactive contact points between neurites, dendritic cells in the spleen and tonsil, etc. (25). The binding of PrPSc with the membrane receptor may change the local environment of the myelin, which may then cause the failing of PrPc glycosylation right after PrPSc anchors PrPc on the membrane. Secondly, non-glycosylated PrPc may be more easily polymerised and transformed into pathogenic PrPSc using PrPSc as the template.

Since configuration shift or transformation of protein may depend on the environmental pH, the configuration of PrPSc formed at pH 4–6 shows stronger resistance to protease than its counterpart formed at pH 7.4. Low pH may stimulate the conversion of the secondary structure from PrPc to PrPSc, and prion protein should be insoluble at pH below 3.5. As a result, configuration changes of PrPSc must be pH-dependent, and tend towards the β-sheet by blocking the digestion sites of the protease at pH below 4.0. The configuration conversion of PrPSc is mainly caused by N-terminal infolding or invagination (9).

It is hardly possible for PrPSc to be internalised after combining with other chaperone molecules as the resulting structures are too large. Perhaps PrPSc could react with different types of trans-membrane molecules on various cell membranes and the resultant reaction between PrPc and PrPSc may convert the normal PrPc into PrPSc (21). When PrPSc appears on the cell myelin sheath, it usually shows decreased membrane tension, buckling, invagination, caveolae, and increased folding and stratification. These could be related to the change in the membrane structure, which is caused by the polymerisation of PrPc and PrPSc complexes. Exogenous PrPSc must be internalised into the cell membrane in order to infect new cells. In this case, PrPSc is able to contact PrPc mainly on the molecular surface at the C-terminal, however, this process is limited by the specificity of prion protein GPI binding to the membrane. Therefore, PrPc may be considered a receptor for the internalisation of PrPSc into the cell. In other words, PrPSc is not able to infect all kinds of animals (24) as the specificity of PrPc involves the species barrier.

Several studies reported that deglycosylated or incompletely glycosylated PrPc is much easier to convert into PrPSc (32). The glycosylation process would be hampered after PrPc binding with PrPSc, and non-glycosylated molecules are prone to accumulate and change configuration, which may also cause mutation in the sequence of PrPc and then transformation to PrPSc (10). Several scientists artificially performed deglycosylation of PrPc and as a result, PrPc accumulated in the cytosol and converted into pathogenic PrPSc. Hence, glycosylation may regulate the configuration changes between PrPc and PrPSc (32). Aggregated particles could also be seen in other experiments as well, but they were not so typical as in our test and so failed to draw the scientists’ attention (19). The characteristic findings of the particle aggregations in the myelin were more obvious and strongly consistent with several other results (20, 26).

However, the multilayer structure and herniation of the myelin formed cannot be explained in the present study. It may be caused by a change in membrane structure or be due to the alteration of polymeric patterns (12).

At day 90, the mice started to show clinical symptoms such as itching and depression. Therefore,
the changes in their brains were observed beginning from 90 dpc, which was a key time point judging from a series of pathological lesions. In the initial stage, the configuration-changed PrPSc in nerve fibres was polymerised as seeds and the misfolded proteins caused configuration disorder. From day 90, similar changes occurred; abnormal PrPSc proteins were polymerised directly and led to the continuous appearance of misfolded proteins. As the proteasomes controlling this kind of misfolded proteins were over-burdened, this led to the formation of abnormal PrPSc protein polymers, the deglycosylated proteins occupying the dominant places, and myelin layering or hyperplasia (25, 28). However, the aggregation of suspicious black particles did not appear, they presented with dissolved or disintegrated forms; we supposed it may be caused by autophagy, proteasome, and inner environmental alterations. These kinds of processes were not really observed very clearly in our experiments, except autophagy that was formed through axoplasm dissected by the cytoplasmic membrane. The polymer of PrPSc with the resistant structures of a protease repressor furnished genuine proof that the normal prion proteins could be changed into abnormal prion proteins. Meanwhile, mixing PrPSc and PrPSc in vitro would start to convert PrPSc to PrPSc under certain conditions (2). The polymerised PrPSc was as toxic to cells as some varieties of similar misfolded proteins, such as membrane-protein-1, presenilin-1, cystic-fibrosis transmembrane conductance regulator (CFTR), and parkin protein. The prion protein was not dissolved but became polymer-like particles in the myelin, while it was soluble in cytoplasm. The damage to the peripheral organelles caused by aggregation of suspicious black particles occurred gradually in the myelin sheath. It may have been the combination of the proliferation of PrPSc with myelin bending, changes of tension, and inner environmental factors which jointly delaminated and aggregated black particles in the myelin sheath. When the layered structure of myelin was completely destroyed along with the gradual homogenisation or dissolution of the suspicious black particle aggregations, pathological lesions would gradually expand to the periphery.

The changes and degeneration between the two myelin sheaths were more violent or faster, which may be due to the fact that more prion proteins were attracted by the prion anchor and more misfolded proteins interacted with each other. In this study, the classical lesions in the nerves were observed, including Wallerian degeneration, myelinat fibre formation, and intramyelinic vacuoles. With the increase of PrPSc in the myelin, the structures of the myelin sheath and plasma membrane were destroyed and then PrPSc diffused into the cells and the surroundings, finally causing the degradation of proteasome and autophagic vacuoles in axoplasm. The organelles may be digested by autophagic vacuoles, leading to death of cells. The degenerated nerve fibres in the experiment had actually lost their normal function, and in clinical examination the mice showed symptoms such as itching (7).

The most obvious pathological lesions observed in mouse brain were the aggregation of particles and stratification in the myelin sheath after infection for 90 days. The emergence of polymeric particles may be caused by PrPSc induction, comprehensive effects of environment within the myelin, and a variety of auxiliary proteins or other molecules. It has been speculated that the conversion of PrPSc to PrPSc was just a configuration change of proteins without the involvement of nucleic acids (14, 30).

As for the pathogenicity of the polymeric protein, orderly aggregated pathogens were highly infectious. The PrPSc in monomeric form was usually harmless, and yet it might be prone to convert into the primary PrPSc aggregates. PrPSc polymers consisted of less than 20 molecules with higher toxicity and faster diffusion ability (2). Thus, prion strain RML should be a highly toxic strain, as no big polymers were observed in the myelin sheath. Toxicity of the low polymerised PrPSc was first manifested in the dysfunctional nerve fibre membrane, extremely swollen myelin, and induction of disintegration and apoptosis of cells. PrPSc is the pathogenic factor of several prions, and its pathogenesis may be different from other prions. The diffusion of polymeric particles has still not been elucidated but may be due to various environmental pH and enzyme activities. The internal environment of the myelin sheath is conducive to the formation of PrPSc and protein misfolding.

PrPSc would eventually reach the brain through either gastrointestinal ingestion or injection into the body, and cause deadly damage to the central nervous system. It was thought that PrPSc was transmitted by the axoplasm of nerve fibres and synapses (2, 28). We speculated that the possible transmission manner of PrPSc was along the myelin sheath by other means than axoplasm. The transmission speed of PrPSc in the axonal cytoplasm membrane cavity is 0.5–3 mm/day (28). Alternatively, the transmission between cells is through endocytosis, and the denatured prion proteins are aggregated into a stable complex. It is gradually transferred along the nerve sheath, possibly through the domino effect (28).

Prion aggregation may be caused by environmental changes, for the cells change their structures in response to specific variations of environmental factors. In this study, the aggregate particles are observed, and we think PrPSc is the causative factor, which has also been found in bacteria of prokaryotic cells (35). So, the structure change of prions may be a universal phenomenon in biology. The previous research proved that the two key positions (23 and 12 bp indels) are sensitive sites where potentially the transformation of PrP into PrPSc could be stimulated (8, 17, 22, 33). In our experiment, we found the myelin started to polymerise from one site and then gradually
spread by domino effect, which may be associated with the excitation sites. Yuan et al. (35) found the structure transformation of prion after inserting a section of prion into the bacterial membrane, while we observed that the transformation might start from myelin sheath in mice. The inability to see the polymeric particles in the cytoplasm may be attributed to the dissolved state.

In summary, electron microscope observation showed widespread presence of particle aggregations and delamination in the axons and dendrites of the myelin sheath in the brain of BALB/c mice injected with pathogenic brain homogenates. Aggregation of particles and stratification occurred much more easily in the adjacent myelin, which demonstrated that the conformational change of prion protein was more likely to occur in the myelin sheath and not in cytoplasm or outside the cells.

Conflict of Interests Statement: The authors declare that there is no conflict of interests regarding the publication of this article.

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