Human prion protein sequence elements impede cross-species chronic wasting disease transmission

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Chronic wasting disease (CWD) is a fatal prion disease of North American deer and elk and poses an unclear risk for transmission to humans. Human exposure to CWD occurs through hunting activities and consumption of venison from prion-infected animals. Although the amino acid residues of the prion protein (PrP) that prevent or permit human CWD infection are unknown, NMR-based structural studies suggest that the β2-α2 loop (residues 165–175) may impact species barriers. Here we sought to define PrP sequence determinants that affect CWD transmission to humans. We engineered transgenic mice that express human PrP with four amino acid substitutions that result in expression of PrP with a β2-α2 loop (residues 165–175) that exactly matches that of elk PrP. Compared with transgenic mice expressing unaltered human PrP, mice expressing the human-elk chimeric PrP were highly susceptible to elk and deer CWD prions but were concurrently less susceptible to human Creutzfeldt-Jakob disease prions. A systematic in vitro survey of amino acid differences between humans and cervids identified two additional residues that impacted CWD conversion of human PrP. This work identifies amino acids that constitute a substantial structural barrier for CWD transmission to humans and helps illuminate the molecular requirements for cross-species prion transmission.

Introduction

Infectious prions cause rapidly progressive, fatal neurodegenerative diseases that can be transmitted zoonotically. For example, bovine spongiform encephalopathy (BSE) prions spread from cattle to humans, causing 229 human deaths from variant Creutzfeldt-Jakob disease (vCJD) as of June 2014 (1–3). In North America, chronic wasting disease (CWD) in deer and elk spreads by horizontal transmission and is remarkably infectious, with a reported incidence of 89% in one captive population (4). Human exposure to CWD prions likely occurs through consumption of venison from CWD-infected deer (5), as both muscle and fat from deer have been shown to contain infectious prions (6, 7). An active surveillance of more than 17,000 US residents revealed that nearly 20% hunt deer or elk, and more than two-thirds have consumed venison (8), yet the risk of developing a prion infection through dietary exposure to CWD remains unclear. Human prion protein-expressing (PrP-expressing) transgenic mice resist CWD infection, suggesting a strong barrier for CWD transmission to humans (9–12); however, certain non-human primates are CWD susceptible (13–16). Similarly conflicting results from studies performed in vitro showed that CWD prions either efficiently or poorly convert human PrP to a pathogenic isoform (14, 17–19). Thus, no consensus has emerged on the susceptibility of humans to CWD prions, and the transmission of CWD to humans remains a major public health concern.

Cross-species prion transmission is influenced by (a) the sequence similarity between the cellular prion protein (PrPc) and the misfolded, aggregated conformer (PrPSc) and (b) the PrPSc conformation (20). PrPSc has an unstructured N terminus and a globular C-terminal domain, arranged in three α-helices and a short anti-parallel β-sheet, together comprising approximately 210 amino acids (21). The tertiary structure of PrPSc is highly conserved among mammals; however, specific amino acid differences between species are hypothesized to impact the intermolecular binding of PrPSc and PrPSc (22). One segment of high sequence diversity is the β2-α2 loop, consisting of residues 165–175 (human numbering) in which only 3 amino acids (P165, Y169, and Q172) are strictly conserved (23, 24).

Nuclear magnetic resonance–based (NMR-based) structural studies suggest that residue differences within the β2-α2 loop and the C terminus would preserve the global shape of PrP but may alter surface hydrogen bonding patterns that influence protein-protein interactions (22). Indeed, β2-α2 loop substitutions were shown to impair prion conversion in vitro (25–28) and in vivo, as transgenic mice that express PrP with 168R or with Y169G, S170N, and N174T substitutions resist infection with mouse-adapted prions (29, 30). Collectively, these studies provide a rationale for investigating the role of the β2-α2 loop in barriers to human infection with CWD.
transmission. Together our results reveal specific amino acids that impair CWD transmission to humans as well as a new determinant for cross-species prion transmission.

Results

Engineering transgenic mice expressing a human-elk chimeric PrP construct. We developed transgenic mice expressing human PrP with four amino acid residues from elk PrP on a Prnp–/– background (Figure 1A), referred to as Tg(HuPrP<sub>166–174</sub>) mice. Transgenic mice expressing human PrP [Tg(HuPrP)] were used as controls (9), and the same plasmid vector was used to generate the two transgenic mouse lines. The Tg(HuPrP<sub>166–174</sub>) mice and the Tg(HuPrP) control mice had comparable PrP C<sub>1</sub> levels in the brain, i.e., approximately 1- to 2-fold higher than those of WT mice (Supplemen-
Figure 2. CWD prions differ biochemically in Tg(HuPrP<sup>Δ166–174</sup>) mice as compared with cervid CWD or human CJD. (A) PrP<sup>Sc</sup> from Tg(HuPrP<sup>Δ166–174</sup>) brain migrates faster and has a glycoform pattern (ratio of di-, mono-, and un-glycosylated PrP<sup>Sc</sup>) different from that of elk CWD by Western blot analysis. Results for six of the seven positive mice are shown here. Red dashes indicate migration of unglycosylated PrP<sup>Sc</sup>. (B) In Tg(HuPrP) mice inoculated with elk CWD, the brain shows no detectable PrP<sup>Sc</sup>, even after NaPTA precipitation. (C) Tg(HuPrP<sup>Δ166–174</sup>) mice infected with deer CWD show a PrP<sup>Sc</sup> electrophoretic migration and glycoform pattern indistinguishable from that of Tg(HuPrP<sup>Δ166–174</sup>) mice inoculated with elk CWD. (D) As in B, no Tg(HuPrP) mice inoculated with deer CWD have detectable PrP<sup>Sc</sup> in the brain. (E) Sub-passaged CWD-Tg(HuPrP<sup>Δ166–174</sup>) (P2) in Tg(HuPrP<sup>Δ166–174</sup>) mice revealed no change in the electrophoretic mobility or glycoform pattern of PrP<sup>Sc</sup> as compared with PrP<sup>Sc</sup> from P1. (F) Comparison of Tg(HuPrP<sup>Δ166–174</sup>)-CWD prions with human sporadic and variant CJD prions. Isolates consisted of type 1 or type 2 sporadic CJD prions from individuals homozygous for methionine (MM) or valine (VV) at PRNP codon 129, Tg CWD, CWD-infected Tg(HuPrP<sup>Δ166–174</sup>). (G) Measurements of di-, mono-, and un-glycosylated PrP<sup>Sc</sup> from CWD-inoculated Tg(HuPrP<sup>Δ166–174</sup>) mice differed significantly from those of CWD-infected elk and deer. **P < 0.01; 2-tailed, unpaired Student’s t test comparing ratio of di- and monoglycosylated PrP (n = 3–6 animals per group).
The Journal of Clinical Investigation

Research Article

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(sCJD) prions, and therefore we inoculated Tg(HuPrPelk166–174) and human PrP may have created a barrier to human sporadic CJD infection.

Tg(HuPrPelk166–174) mice was processed similarly to WT PrP, as both were glycosylated and anchored in lipid rafts together with flotillin (Supplemental Figure 1B). Since certain transgenic mice expressing mutant PrP develop spontaneous prion disease, we examined 29 aged Tg(HuPrP166–174) mice (350–676 days old), yet found no evidence of prion disease. Mice had no evidence of neurological impairment, PrP deposits on histologic sections, or PrP aggregates detected by biochemical assays (Supplemental Figure 1C).

Tg(HuPrPelk166–174) mice develop CWD prion infection. We inoculated Tg(HuPrP166–174) and Tg(HuPrP) mice with CWD prions from a naturally infected elk or with uninfected cervid brain (mock control). Animals were examined every other day for behavioral changes or neurological impairment. None of the Tg(HuPrP) mice inoculated with elk prions developed clinical disease by ≥587 days after inoculation (n = 12) (Figure 1, B and C), consistent with previous reports (9). Three of 12 mock-inoculated Tg(HuPrPelk166–174) mice died of non-prion-related causes. In contrast, 7 of 8 (88%) Tg(HuPrP166–174) mice inoculated with elk CWD prions manifested terminal signs of neurologic disease, including immobility, progressive weight loss, hind leg clasp, and disorientation (268 ± 16 days post-inoculation [dpi], mean ± SEM) (Figure 1, B and C, and Supplemental Videos). Tg(HuPrP166–174) mice were also susceptible to mule deer prions (3 of 4 mice infected, incubation period 271 ± 54 dpi).

Tg(HuPrPelk166–174) mice show a delay in developing human prion infection. We reasoned that the new elk β2-α2 loop sequence in human PrP may have created a barrier to human sporadic CJD (sCJD) prions, and therefore we inoculated Tg(HuPrPelk166–174) and Tg(HuPrP) mice with sCJD. Subsequently, all Tg(HuPrP) mice developed terminal prion disease by 177 ± 3 dpi (n = 6), whereas all 3 Tg(HuPrPelk166–174) mice developed terminal prion disease by 280 ± 10 dpi, an approximately 60% lengthening of the incubation period (Figure 1, E and F, and Supplemental Figure 2). These findings indicate that elk residues at positions 166, 168, 170, and 174 of human PrP enable CWD-induced conversion yet delay the development of terminal CJD infection.

Thalamic and hypothalamic prion deposits in CWD-infected Tg(HuPrPelk166–174) mice. Neuropathologic assessment of brains from CWD-inoculated Tg(HuPrPelk166–174) mice showed lesions typical of prion disease, including focal spongiform change, gliosis, and diffuse and punctate PrPSc deposits in the thalamus and hypothalamus (Figure 1D). Lesions were restricted primarily to the thalamus and hypothalamus, whereas the Tg(CerPrP) mice showed severe, widespread spongiform changes in the cerebral cortex, hippocampus (dentate gyrus), and putamen, with severe neuronal loss in the cerebellum (9).

The proteinase K-resistant (PK-resistant) core of PrP, a hallmark of prion infection, was detected biochemically in all elk CWD–inoculated Tg(HuPrPelk166–174) mice, with terminal disease (n = 7) but in none of the Tg(HuPrP) mice (Figure 2, A and B). Inoculation with deer CWD produced similar results in Tg(HuPrPelk166–174) and Tg(HuPrP) mice (Figure 2, C and D). Thus, the CWD-inoculated Tg(HuPrPelk166–174) mice presented the classic features of prion disease in the brain, both histopathologically and biochemically. HuPrP166–174-CWD prions from one infected mouse were inoculated into Tg(HuPrP) and Tg(HuPrPelk166–174) mice. No clinical or biochemical evidence of prion disease was detected in any Tg(HuPrP) mice by 545 dpi (n = 5), whereas all Tg(HuPrPelk166–174) mice developed terminal prion disease by 203 ± 19 dpi (n = 4), indicating that a substantial barrier exists between the two groups despite the fact that they differ by only 4 residues.

Human-elk chimeric prions show altered biochemical properties as compared with elk CWD prions. The PrPSc conformation may be preserved or changed profoundly when transmitted to a species with a different PrP sequence (31). We therefore compared the biochemical properties of HuPrPelk166–174-CWD with those of elk CWD prions and found consistent evidence supporting the former’s unique conformation. First, the electrophoretic mobility of PK-resistant HuPrPelk166–174-CWD increased, indicating a smaller PK-resistant core (Figure 2, A and C), which also differed from human sporadic and variant CJD (Figure 2F). Second, in HuPrPelk166–174 prions, we noted a clear predominance of diglycosylated PrPSc, whereas elk prions contained nearly equal proportions of di- and monoglycosylated PrPsc (60:20:20 versus 40:40:20, di-, mono-, and unglycosylated PrPSc, respectively) (Figure 2G). Third, HuPrPelk166–174 prions were consistently more resistant to guanidine HCl-induced (Gdn-HCl–induced) unfolding (32) than elk CWD (Figure 3, A–C). Last, the relative levels of PK-resistant PrPsc differed; HuPrPelk166–174 prions were composed of approximately two-thirds PK-resistant PrPsc (66% ± 5%; mean ± SEM), whereas elk contained 3-fold less PK-resistant PrPsc (20% ± 3%) (Figure 3, D and E, and Supplemental Figure 3). These four independent measures indicated that elk and HuPrPelk166–174 CWD vary biochemically, consistent with a change in the CWD conformation upon conversion to the human-elk chimeric sequence. The elk residues also modified the CJD glycoform profile (Figure 3, F and G), an outcome that lends further support to the concept that the β2-α2 loop region impacts PrPSc–PrPPrSc interactions.

We next investigated adaptation of the new human-elk chimeric prions by further passaging in Tg(HuPrPelk166–174) and Tg(HuPrP) mice. Sub-passage of HuPrPelk166–174 prions in Tg(HuPrPelk166–174) mice revealed no significant change in the electrophoretic mobility, glycoform pattern (Figure 2E), or PrPsc deposition sites (Supplemental Figure 4), indicating that the dominant conformation was preserved. Notably, the incubation period decreased significantly (24%, P < 0.05, Student’s t test) upon second passage. To assess whether these findings could be reproduced in vitro in a conversion assay, we performed protein misfolding cyclic amplification (PMCA) using brains from the Tg(HuPrPelk166–174) and Tg(HuPrP) mice as PrPSc substrates. Indeed, the HuPrPelk166–174 was converted by CWD within five rounds, whereas the HuPrP was not converted by CWD, even after 10 rounds of amplification (Table 1 and Supplemental Figure 5).
168Q and 170N substitutions markedly reduce the barrier for CWD conversion of human PrP. To address how each of the four residue substitutions impacts conversion by CWD prions, we generated human PrP \(^\text{C}\) having single elk substitutions at positions 166, 168, 170, or 174 in a PrP\(^\text{C}\)-deficient RK13 cell line. Cell lysates containing the human PrP\(^\text{C}\) variants or cervid PrP\(^\text{C}\) were seeded with elk CWD prions and subjected to repeated cycles of sonication for 24 hours in a cell lysate–based PMCA assay (26, 33, 34), which previously reproduced in vivo prion susceptibility (28).

Here, we found that HuPrP with either the E168Q or S170N substitution was converted by CWD, albeit at low levels (17%–26%) relative to conversion of cervid PrP\(^\text{C}\) (Figure 4, A and B). We next tested HuPrP with elk substitutions at 168 and 170 together, and found remarkably robust conversion equivalent to that of cervid PrP\(^\text{C}\) (Figure 4, C and D). Intriguingly, HuPrP with elk substitutions at positions 166, 168, 170, and 174 resulted in only approximately 10% conversion, revealing more efficient conversion in HuPrP with two versus four elk substitutions (Figure 4, C and D).

Since the efficiency of cross-species prion conversion depends on amino acid sequence similarity, the less-efficient conversion with four versus two residue substitutions was surprising and unexpected. As residue 166 interacts with the C-terminal segment of helix \(\alpha_3\) (residues 218–226) (35), we tested whether a long-range interaction between residues 166 and 219, where human and cervid PrP differ, could explain the low conversion efficiency. Thus, we next measured conversion of HuPrP with elk residues at 166, 168, 170, 174, as well as 219, yet found only a negligible increase in conversion with the addition of the 219 substitution (Figure 4, C and D), suggesting that the previous results were due to local effects within the \(\beta_2-\alpha_2\) loop. Further testing of HuPrP with elk residues at 166, 168, and 170, or at 168, 170, and 174, revealed that the N174T elk substitution was impairing CWD-induced conversion of human-elk chimeric PrP\(^\text{C}\) (Figure 4, C and D).

We noted that together, the E168Q and S170N elk substitutions in HuPrP created an asparagine- and glutamine-rich, highly aggregation-prone \(\beta_2-\alpha_2\) loop sequence (QYNNQNNF, from 168 to 175) identical to that of bank vole, a species known to be remarkably susceptible to prions from many different species (36). Thus, these data collectively suggest that a highly aggregation-prone \(\beta_2-\alpha_2\) loop segment can override PrP\(^\text{Sc}\)-PrP\(^\text{C}\) amino acid sequence mismatches. We propose that prion conversion between two species is determined not only by (a) PrP amino acid sequence similarity and (b) PrP\(^\text{Sc}\) conformation, but also (c) the amyloid-forming propensity of key segments of the host PrP\(^\text{C}\) protein.

Positions 143 and 155 also impact the CWD species barrier. To identify other amino acids that impact CWD-seeded conversion of human PrP\(^\text{C}\), we generated human PrP\(^\text{C}\) having the single elk substitutions I138L, S143N, H155Y, I184V, and V203I. At each cervid polymorphic site, the human PrP sequence matched the cervid
Table 1. Summary of CWD- and CJD-seeded conversion of HuPrPelk166–174 and HuPrP by PMCA

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*Representative results of two experiments in which samples were seeded with either of two CWD isolates (CWD1 and CWD2) or the MM1 subtype of CJD. The number of positive samples (bold) is shown as a fraction of the total replicates per experiment.

Discussion

In 1996, ten cases of new variant CJD were reported in the UK with suspected links to the BSE epidemic (37, 38), and extensive evidence now supports that cattle BSE crossed the species barrier and infected humans (39). With the realization that animal prions could transmit to humans, concerns arose that CWD in cervids may lead to cases of a novel form of CJD. Nevertheless, four laboratories have independently reported that transgenic mice expressing human PrP resist CWD infection, suggestive of a strong barrier to infection (9–12). Here we have identified the specific residues in human PrP that modulate CWD transmission. We report that CWD transmits to mice expressing a human-elk chimeric PrP and show that the CWD-human species barrier is largely maintained by the human-specific amino acids within the β2-α2 loop. Within the loop, human residues E168 and S170 are significant inhibitors of CWD conversion, as evidenced by in vitro conversion experiments. Human residues S143 and H155 also contribute to the CWD to human transmission barrier.

Elucidating the determinants of cross-species prion transmission. Prions can transmit between different species; however, infection of a new species is typically characterized by prolonged, variable incubation periods and low attack rates (41, 42). Known determinants of interspecies prion conversion are (a) PrP\(^{c}\) and PrP\(^{Sc}\) sequence similarity and (b) the conformation of the infectious prion (43, 44). Here we propose to add a third determinant, the presence of glutamines or asparagines in host PrP\(^{c}\) within key interaction segments. These key segments will vary depending upon exposure in the PrP\(^{c}\) conformation. We found that human PrP with the E168Q and S170N substitutions was readily converted by CWD, whereas adding the N174T substitution, which increases homology with elk CWD but removes an asparagine, paradoxically led to a massive decrease in conversion. These results suggest that for CWD, the β2-α2 segment is accessible and interacts with host PrP\(^{c}\). These data also indicate that the presence of N/Qs in the loop can be a strong determinant for conversion that overrides certain sequence differences.

Glutamine- and asparagine-rich protein segments are proposed to play a role in protein aggregation due to side chain hydrogen bonding among amide groups that stabilizes adjacent β-strands (45–47). Asparagine residues are prevalent in the PrP\(^{c}\) of bank voles, a species that is highly susceptible to CWD, sheep scrapie, and CJD, despite having many PrP\(^{c}\) sequence mismatches with these infectious prions (48–51). Indeed, bank voles were recently designated the “universal acceptor for prions” (36). Similar to the bank vole, hamsters are susceptible to CWD and other prions (42, 52–54) and also have an N/Q-rich loop. Thus, it is possible that the bank vole-like sequence in the human β2-α2 loop creates a permissive host PrP\(^{c}\) sequence that is converted by prions from other species, despite sequence mismatches. Consistent with the observation that N/Q residues lead to a lower species barrier, fewer N/Qs in key segments may be protective, as polymorphisms that replace a single N/Q residue — for example, Q168R in sheep PrP\(^{c}\) (55, 56), Q219K in mouse PrP\(^{c}\) (30), and Q219K in goat PrP\(^{c}\) (57) (all human numbering) — correlate with profound resistance to certain prions in vivo.

Accumulating evidence suggests that the β2-α2 loop governs certain prion transmission barriers and conformational properties, reminiscent of Sup35 yeast prion segments, also shown to govern species barriers and strains (58). In transgenic mice, S170N and N174T loop substitutions increased susceptibility to CWD and hamster prions, led to resistance to sheep or cattle prions, and altered the RML mouse prion conformation (59). Here we show that replacing four human amino acids in the β2-α2 loop enabled infection with elk CWD and created a partial barrier for human CJD, as evidenced by the delayed infection kinetics. Additionally, HuPrPelk166–174-CJD showed a different glycosylation pattern as compared with human CJD prions, suggesting the emergence of a new CJD conformation. Surprisingly, passage of HuPrPelk166–174-CWD prions revealed efficient infection of all Tg(HuPrPelk166–174) mice, but not Tg(HuPrP) mice, despite only four amino acid differences in PrP.
A proposed structural mechanism for the CWD-to-human transmission barrier. How do short PrP segments control prion conversion? Microcrystal structures of segments from amyloid-like fibers invariably reveal pairs of tightly packed β-sheets, in which complementary side chains interdigitate in a dry “steric zipper” interface (60). We hypothesize that efficient prion conversion requires donor and recipient PrP loop segments to form a tight steric zipper, whereas side chain mismatches lead to steric clashes and cavities, prevent conversion, and may account for species barriers in prion disease (61, 62). Consistent with this hypothesis, microcrystal structures of human and cervid β2-α2 loop segments belong to different classes of steric zippers (60, 62), and our computational analysis suggests that the human and cervid loop segment do not form complementary steric zippers (Figure 5, Supplemental Figure 6, and Supplemental Table 1). We found that the S170 (human) and N170 (elk) side chains leave a cavity within the zipper core due to poor packing (Figure 5 and Supplemental Figure 6). Also, the E168 (human) and Q168 (elk) residues steric clash, which hinders tight, highly complementary side chain interactions between β-sheets in the zipper models (Figure 5 and Supplemental Figure 6). In summary, the zipper models suggest that residue mismatches between human and elk PrP at positions 168 and 170 would not support efficient conversion and may explain our results at the atomic level.

In conclusion, we have identified the β2-α2 loop sequence of human PrP as a major barrier to PrP conversion by CWD prions. The human-specific residues in the β2-α2 loop, particularly E168 and S170, appear to substantially raise the energetic barrier for conversion of human PrP by CWD prions.
ground. Zygosity of the Tg(HuPrP<sup>166–174</sup>) mice was determined by TaqMan qPCR performed at the UC Davis Mouse Biology Program. PrP expression in the brains of transgenic and WT mice was assessed by Western blot analysis using the anti-PrP monoclonal antibody POM19 (provided by Adriano Aguzzi, University Hospital of Zurich, Zurich, Switzerland) (66). Mice were maintained under specific pathogen–free conditions.

**Prion inoculations.** WT (FVB), Tg(HuPrP<sup>166–174</sup>), or Tg(HuPrP<sup>166–174</sup>) (homozygous mice generated from the previously reported Tg40) transgenic mice (groups of <i>n</i> = 4–10 mice) were intracerebrally inoculated into the left parietal cortex with 20–30 μl of brain homogenate (1%–5%) containing CWD prions from a naturally infected elk or mule deer previously shown to contain infectious prions (67) or from a human with sporadic CJD. Uninfected brain homogenates were inoculated into mice of the same genotypes as negative controls. Mice were monitored three times weekly, and TSE was diagnosed according to clinical criteria including ataxia, kyphosis, stiff tail, hind leg clasp, and hind leg paresis. Mice were sacrificed at the onset of terminal neurological disease when showing such signs as weight loss, tremors, slow movements, and severe kyphosis, or at approximately 600 dpi. Incubation period was calculated as the day of inoculation to the day of terminal clinical disease.

**Methods**

**Generation of transgenic mice.** Development of the Tg(HuPrP)-expressing mice was previously described (9). Constructs for the HuPrP<sup>166–174</sup> transgenes were based on the same human PrP in the half-genomic plasmid (pHGPRP) used for the Tg(HuPrP) mice (9, 65). Site-directed mutagenesis was performed to replace the sequence coding for 166, 168, 170, and 174. The final construct was confirmed by sequencing. One error-free clone was chosen for the creation of transgenic mice by pronuclear microinjection, performed at the UC San Diego Transgenic Mouse Core facility. Founder pups were screened by tail DNA PCR using the following primers: forward, 5′-CAACCGAGCT-GAAGCATTC; reverse, 5′-CCCTCTCGTACTGGGTGATAC. All founder mice that carried the transgene were bred with FVB/Prnp<sup>0/0</sup> mice (provided by Stanley Prusiner’s laboratory, UCSF, San Francisco, California, USA) to achieve transgenic mice with a Prnp<sup>0/0</sup> background.
kosił and digested with an endonuclease (Benzonase, Sigma-Aldrich), followed by treatment with 20 μg/ml proteinase K at 37°C for 30 minutes. After the addition of NaPTA, MgCl₂, and protease inhibitors (Complete-TM, Roche), extracts were incubated at 37°C for 30 minutes and centrifuged at 18,000 g for 30 minutes at 37°C. Pellets were resuspended in LDS loading buffer (Invitrogen), and samples were heated to 95°C prior to electrophoresis through a 10% Bis-Tris gel (Invitrogen) and transferred to a nitrocellulose membrane by wet blotting. Proteins were detected with anti-PrP antibodies 3F4 (MAB1562, Millipore) and POM19 (66), followed by an HRP-conjugated anti-mouse IgG antibody (Jackson ImmunoResearch Laboratories Inc.). Signals were visualized using a chemiluminescent substrate (Super signal West Dura, Thermo Scientific) and an LAS-4000 imager (Fuji-film). Prion-infected or uninfected brain samples were also subjected to NaPTA precipitation and blotting to serve as controls.

**Histopathology and immunohistochemistry.** Five-micrometer-thick sections were cut onto positively charged glass slides and stained with hematoxylin and eosin or immunostained using antibodies for glial fibrillary acidic protein (GFAP). For GFAP staining, sections were deparaffinized, rehydrated, and quenched with 3% hydrogen peroxide for 10 minutes. Sections were then incubated in 20 μg/ml PK for 10 minutes, blocked, and incubated with anti-GFAP monoclonal antibody (M0761, Dako) for 45 minutes, followed by biotinylated anti-mouse IgG (Jackson ImmunoResearch Laboratories Inc.) for 30 minutes, streptavidin-HRP (Jackson ImmunoResearch Laboratories Inc.) for 30 minutes, and DAB substrate for 7 minutes. Sections were counterstained with hematoxylin.

**Paraffin-embedded tissue blots.** Five-micrometer-thick sections were collected onto 0.45-μm nitrocellulose membranes (Bio-Rad) and dried at room temperature overnight and then at 55°C for 30 minutes. Membranes were then incubated in xylene and serially rehydrated in 100% isopropanol, 70% isopropanol, and distilled water with 0.1% Tween-20 for 10 minutes each. To improve tissue adherence, membranes were dried. After a brief rinse with TBST (10 mM Tris-HCl [pH 7.8], 100 mM NaCl, 0.05% Tween-20), membranes were incubated in 50 μg/ml PK for 10 minutes, blocked, and incubated with anti-PrP monoclonal antibody (M0761, Dako) for 45 minutes, followed by biotinylated anti-mouse IgG (Jackson ImmunoResearch Laboratories Inc.) for 30 minutes, streptavidin-HRP (Jackson ImmunoResearch Laboratories Inc.) for 30 minutes, and DAB substrate for 7 minutes. Sections were counterstained with hematoxylin.

**Cell lysate-based PMCA.** Cervid Prnp (G96, M132, S225, Q226 [cervid numbering]) and human PRNP (M129) in the pcDNA3.1C vector (Invitrogen) containing the 3F4 epitope tag (109M, I12M) were generated by site-directed mutagenesis as previously described (28). PrPc contained the highly conserved amino-terminal coding sequence (residues 23–111) of cervid Prnp and the cervid alanine residue at position 230. Site-directed mutagenesis (Agilent) was used to generate an array of cervid Prnp and human PRNP plasmids, each with amino acid substitutions as described in Results. Human PRNP with the M112V was not generated, as this would have removed the 3F4 epitope used for detection. The final constructs were confirmed by sequencing.

**Velocity gradient sedimentation.** 10% brain homogenates prepared in PBS containing 9% sucrose and protease inhibitors (CompleteMini, Sigma-Aldrich) were centrifuged at 750 g for 5 minutes at 4°C. Supernatant was mixed with an equal volume of 2% Triton X-100 in PBS containing 9% sucrose. Samples were brought to a final concentration of 40% OptiPrep (Sigma-Aldrich) using flotation buffer, then overlaid with 30% and 50% OptiPrep prepared using flotation buffer (10 mM MES containing 1% Triton X-100, 2 mM EDTA, 1 mM DTT, and protease inhibitors). All steps were performed on ice. Samples were centrifuged at 200,000 g for 18 hours at 4°C using an MLA-80 rotor (Beckman-Coulter). Fractions 1–13 were collected from top to bottom.
Size exclusion chromatography of PrP\textsuperscript{Sc}. CWD-infected elk and CWD-infected Tg(HuPrP\textsuperscript{PrP\textsubscript{27–132}}) mouse brain homogenates were digested using Benzonase (Millipore) with 50 mM MgCl\textsubscript{2} for 20 minutes at 37°C, mixed with 1% sarkosyl for 20 minutes at 37°C, and centrifuged at 2,000 g for 5 minutes. Supernatants were injected onto a Fast Performance Liquid Chromatography (FPLC) column (Superose 6, 3.2/30, GE Healthcare Life Sciences) using running buffer (50 mM sodium acetate [pH 8.5] containing 0.1% sarkosyl) at a flow rate of 60 μl/min, and 24 fractions were collected. Fractions 2–23 from CWD-infected or uninfected elk and CWD-infected or uninfected Tg(HuPrP\textsuperscript{PrP\textsubscript{27–132}}) mice were subjected to SDS-PAGE, followed by Western blotting using anti-PrP antibodies POM19 (elk) and 3F4 (mice).

15B3 immunoprecipitation for PrP\textsuperscript{Sc}-specific detection. To semiquantitatively measure PrP\textsuperscript{Sc} in FPLC fractions, PrP\textsuperscript{Sc}- and PrP\textsuperscript{Sc}-containing fractions (fractions 1–7 and fractions 8–24, respectively) were each pooled and analyzed for PrP\textsuperscript{Sc}. Pooled fractions were incubated in PBS and protease inhibitors (Roche), 0.1% sarkosyl, and rat anti-mouse Dynabeads (Invitrogen) overnight at 22°C with rotation to remove proteins that bind to unlabeled beads. Beads were removed, and each sample was incubated with anti-PrP antibody 15B3-conjugated Dynabeads for 3 hours at 22°C with rotation. The beads were washed four times with wash buffer (10 mM Tris-HCl [pH 7.4] containing 1% Triton X-100, 150 mM NaCl, and 5 mM EDTA) and boiled in loading dye for 5 minutes prior to SDS-PAGE and detection using biotinylated monoclonal antibody POM1 (66).

Computational analysis of zipper models for PrP conversion. To understand the molecular mechanism of cross-species prion transmission, we sought structural models of elk and human PrP side chains within the β2–α2 loop to explain the affects of different amino acid substitutions on CWD conversion in vitro. The modeling strategy was similar to one that was previously described (28). The zipper structure of PrP peptides was built using Rosetta software (https://www.rosetta-commons.org/). The segments were modeled as a parallel β-sheet by using the template backbone structures of both the elk prion NQNTF (PDB code 3FVA; ref. 70) and the yeast prion Sup35 GNNQQQNY zipper (PDB code 1YPJ; ref. 71). The pair of β-sheets was assembled by exploring all four possible arrangements (class 1–3; the class 1 zipper structure has 2 arrangements) (60). The zipper structure of PrP peptides was then refined by simultaneously optimizing the rigid-body degree of freedom between the β-sheets, side chain and backbone torsions of each β-strand, guided by full-atom Rosetta energy functions. Taking advantage of the recently developed symmetry implementation in Rosetta, the fibril symmetry of each peptide subunit is restrained to assure that symmetrical geometry is satisfied during the whole optimization process. Finally, the models were ranked based on Rosetta full-atom energy (the sum of Lennard-Jones potential for non-bond interactions, implicit solvation energy, hydrogen bond energy, and others) and the packing (shape complementary score and buried solvent-accessible surface area) between β-sheets. Two final zipper models with strongest predicted Rosetta energy and best shape complementary score were selected (classes 1 and 3).

For each selected final structure (classes 1 and 3), the amino acid substitutions were introduced by RosettaDesign. The new models for each substitution were optimized using the same refinement procedure. The Rosetta full-atom energy and the packing between β-sheets for different substitutions were calculated. The analysis and scores of the class 3 zipper structure are summarized in Supplemental Table 1.

Statistics. P values of less than 0.05 were considered statistically significant. The incubation periods for groups of prion-infected mice were compared by log-rank (Mantel-Cox) test. The average incubation period upon first and second passage of elk CWD in Tg(HuPrP\textsuperscript{PrP\textsubscript{27–132}}) mice was additionally compared with a 2-tailed Student’s t test. The ratios of di- and monoglycosylated PrP\textsuperscript{Sc} and PrP\textsuperscript{Sc} stability in prion-infected brains were assessed by 2-tailed unpaired Student’s t tests. Results from in vitro conversion experiments (n = 3–6 experimental replicates) using human PrP variants were compared by 2-tailed unpaired Student’s t test. All data are presented as mean ± SEM unless otherwise noted.

Study approval. All procedures involving live animals were approved by the UC San Diego IACUC (protocol number S08037).

Acknowledgments

We thank Michael Oldstone, Edward Hoover, and Steven Gonias for helpful discussions; Gary Landucci, Donald Forthal, and Alan Barbour for biosafety support and discussions; Tran Phan for excellent technical support; and the animal caretakers at UCSD and UC Irvine. We thank Jean Jewell for CWD inocula used in the PMCA experiments. This study was supported by the NIH (NS055116, NS069566, NS076896, and U54AI0359 [to C.J. Sigurdson] and AG029430 [to D.S. Eisenberg]), national grants from Spain (AGL2012-37988-C04-01, CTP2013-P05, and EFA205/11 [to J. Castilla]) and a fellowship from the Morris Animal Foundation (D13ZO-419 to T.D. Kurt).

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