Epigenetic Dominance of Prion Conformers

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Epigenetic Dominance of Prion Conformers

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Abstract

Although they share certain biological properties with nucleic acid based infectious agents, prions, the causative agents of invariably fatal, transmissible neurodegenerative disorders such as bovine spongiform encephalopathy, sheep scrapie, and human Creutzfeldt Jakob disease, propagate by conformational templating of host encoded proteins. Once thought to be unique to these diseases, this mechanism is now recognized as a ubiquitous means of information transfer in biological systems, including other protein misfolding disorders such as those causing Alzheimer’s and Parkinson’s diseases. To address the poorly understood mechanism by which host prion protein (PrP) primary structures interact with distinct prion conformations to influence pathogenesis, we produced transgenic (Tg) mice expressing different sheep scrapie susceptibility alleles, varying only at a single amino acid at PrP residue 136. Tg mice expressing ovine PrP with alanine (A) at (OvPrP-A136) infected with SSBP/1 scrapie prions propagated a relatively stable (S) prion conformation, which accumulated as punctate aggregates in the brain, and produced prolonged incubation times. In contrast, Tg mice expressing OvPrP with valine (V) at 136 (OvPrP-V136) infected with the same prions developed disease rapidly, and the converted prion was comprised of an unstable (U), diffusely distributed conformer. Infected Tg mice co-expressing both alleles manifested properties consistent with the U conformer, suggesting a dominant effect resulting from exclusive conversion of OvPrP-V136 but not OvPrP-A136. Surprisingly, however, studies with monoclonal antibody (mAb) PRC5, which discriminates OvPrP-A136 from OvPrP-V136, revealed substantial conversion of OvPrP-A136. Moreover, the resulting OvPrP-A136 prion acquired the characteristics of the U conformer. These results, substantiated by in vitro analyses, indicated that co-expression of OvPrP-V136 altered the conversion potential of OvPrP-A136 from the S to the otherwise unfavorable U conformer. This epigenetic mechanism thus expands the range of selectable conformations that can be adopted by PrP, and therefore the variety of options for strain propagation.

Introduction

Prion-mediated phenotypes and diseases result from the conformationally protein characteristics of particular amyloidogenic proteins. The prion state has the property of interacting with proteins in their non-prion conformation, thus inducing further prion conversion. The prion phenomenon has been described for a variety of different proteins involved in diverse biological processes ranging from translation termination in yeast, memory in Aplysia, antiviral innate immune responses [1], and most recently the action of the p53 tumor suppressor [2]. Since the prion and non-prion conformations have differing biological properties, the net result of this replicative process is protein-mediated information transfer, the characteristics of which vary from prion to prion. The ubiquity of prion replication indicates that this is a wide-ranging of means of information transfer in biological systems.

In the case of mammalian neurodegenerative diseases the prion state is pathogenic as well as transmissible. A hallmark of such conditions is the inexorable progression of pathology between synaptically connected regions of the central nervous system (CNS), consistent with advancing cell-to-cell prion spread. Experimental transmission in several settings has been convincingly demonstrated in the case of the amyloid beta (Aβ) peptide which features prominently in Alzheimer’s disease (AD), the intracytoplasmic protein tau, also involved in AD as well as various neurodegenerative diseases referred to as taopathies, and α-synuclein, the primary constituent of Lewy bodies found in Parkinson’s disease (PD) [1,3].

The proteotypic and best-characterized prion diseases are the transmissible spongiform encephalopathies (TSEs) of animals and humans, including sheep scrapie, bovine spongiform encephalopathy (BSE), chronic wasting disease (CWD) of cervids, and human Creutzfeldt-Jakob disease (CJD). TSEs result from conformational
Prions are infectious proteins, originally discovered as the cause of a group of transmissible, fatal mammalian neurodegenerative diseases. Propagation results from conversion of the host-encoded cellular form of the prion protein, PrP^C, to the corresponding prion, or scrapie form, PrP^sc. Since TSEs share numerous properties with nucleic acid-based pathogens, including agent host-range, stable strain properties, and the ability to mutate and respond to selective pressure, early researchers assumed a viral etiology for these diseases. While this is not the case, the unequivocal infectivity of TSEs set these prions apart. Their singular capacity to cause fatal neurodegeneration in genetically tractable animal models, and the ability to propagate and quantify infectivity, in vivo, in cell culture or cell-free conditions, provide unparalleled settings to elucidate general mechanisms and devise integrated therapeutic approaches for all diseases involving conformational templating [4].

TSEs have long incubation periods ranging from months to years, are invariably fatal, and currently incurable. While a variant of CJD (vCJD) is unequivocally linked to prions causing BSE [5], the zoonotic potential of other TSE’s remains uncertain. Whereas all TSEs, including human genetic and sporadic forms, are neurodegenerative diseases involving conformational templating [4].

The influence of the OvPrP A/V136 dimorphism on scrapie strain-specific pathogenesis

We created Tg mice expressing OvPrP encoding either A or V at residue 136. Using semi-quantitative Western and immuno dot blotting we ascertained that levels of expression in the CNS of Tg(OvPrP-A136)3533/− mice and Tg(OvPrP-V136)4166/− mice were close to that of PrP expressed in the CNS of wild type mice (Fig. 1A).

Both lines of Tg mice tolerated these levels of expression without spontaneously developing recognizable signs of disease (Table 1). In contrast, Tg mice of both genotypes intracerebrally inoculated with brain homogenates from scrapie-affected sheep succumbed to the neurological effects of prion disease following variable incubation periods (Table 1). Rapid disease onset occurred following inoculation of Tg(OvPrP-V136)4166/− mice with SSBP/1 prions [15,16], which consistently produced an ~130 d mean incubation time. While SSBP/1 also caused disease in Tg(OvPrP-A136)3533/− mice, mean incubation times were ~230 to 280d longer (Fig. 1B and Table 1). In contrast, CH1641 prions [17] induced disease in all inoculated Tg(OvPrP-A136)3533/− mice with a mean ~310 d onset of disease, whereas no disease was registered in Tg(OvPrP-V136)4166/− mice after >560d. These distinct transmission profiles are consistent with previously recognized strain differences between SSBP/1 and CH1641 scrapie prions [17]. Consistent with this notion, western blot analysis of proteinase K-treated brain extracts of diseased Tg(OvPrP-A136)3533/− mice confirmed that the molecular profiles which distinguish PrP^sc constituting SSBP/1 and CH1641 prions [18] were maintained upon transmission (Fig. 1C).

These results demonstrate that Tg(OvPrP-A136)3533/− and Tg(OvPrP-V136)4166/− mice are capable of distinguishing scrapie strain-specific transmission patterns, and in turn that these properties are influenced by the A/V136 dimorphism.

The influence of the OvPrP A/V136 dimorphism on PrP^sc conversion kinetics, conformation, and CNS deposition

Previous studies revealed a positive correlation between PrP^sc conformational stability and the incubation times of mouse and cervid prions [19,20], but not of hamster prions [21,22]. We performed guanidine denaturation treatments on PrP^sc in brain extracts of SSBP/1 infected Tg(OvPrP-V136)4166/− mice with short incubation times and SSBP/1 infected Tg(OvPrP-A136)3533/− mice with long incubation times. Analyses using mAbs revealed distinct stability curves for OvPrP^sc-V136 and OvPrP^sc-A136. The conformational stability of OvPrP^sc-V136 was lower than OvPrP^sc-A136 in the range of GdnHCl concentrations between 1 and 2 M, Fig. 2A and GdnHCl_1/2 values were 1.78 and 2.17 respectively. This confirmed that the
conformation of OvPrPSc-V136 produced in Tg(OvPrP-V136)4166/+ mice with rapid incubation times was less stable than OvPrPSc-A136 produced in Tg(OvPrP-A136)3533/+ mice with longer incubation times. We refer to these conformers as unstable (U) and stable (S), and to the rapidly and slowly propagating prions composed of these conformers as SSBP/1-V136(U), and SSBP/1-A136(S).

We then used histoblotting [23], a widely used method for characterizing strain-specific differences in PrP Sc distribution [20,24], with mAb 6H4 to characterize OvPrP Sc-A136(S) and

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**Table 1. Prion disease in transgenic mice expressing different ovine PrP scrapie susceptibility alleles.**

<table>
<thead>
<tr>
<th>Isolate</th>
<th>Genotype</th>
<th>Mean time to disease onset ± standard error of the mean (SEM), days¹</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Tg(OvPrP-V136)14166²</td>
<td>Tg(OvPrP-A136)3533¹⁵</td>
</tr>
<tr>
<td>None</td>
<td>&gt;552 (0/4)</td>
<td>&gt;429 (0/5)</td>
</tr>
<tr>
<td>SSBP/1</td>
<td>A136/V136</td>
<td>132±2 (8/8)</td>
</tr>
<tr>
<td>SSBP/1</td>
<td>A136/V136</td>
<td>133±1 (15/15)</td>
</tr>
<tr>
<td>CH1641</td>
<td>A136/V136</td>
<td>&gt;564 (0/6)</td>
</tr>
</tbody>
</table>

¹The number of mice developing clinical signs of prion disease divided by the original number of inoculated mice is shown in parentheses.

²Inoculations performed in Telling lab.

³Inoculations performed in Hunter lab.

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OvPrPSc-V136(U) deposition in the CNS. While OvPrPSc-A136(S) had a punctate pattern of accumulation throughout the midbrain, pons, and oblongata of slow incubation time Tg(OvPrP-A136)3533/+ mice (Fig. 3A), the neuroanatomical distribution of OvPrPSc-V136 in the same sections of rapid incubation time Tg(OvPrP-V136)4166/+ mice was distinctly different, being more intense and diffusely deposited than OvPrPSc-A136(S) (Fig. 3B). Since Tg(OvPrP-A136)3533/+ and Tg(OvPrP-V136)4166/+ mice were both engineered using the same cosSHa.Tet cosmid vector which drives expression from the PrP gene promoter, we conclude that these differences are not the result of expression OvPrP-A136 and OvPrP-V136 in different neuronal populations.

The OvPrP A/V136 dimorphism influences strain-specific PrP conversion in vitro

We used brain extracts of Tg(OvPrP-A136)3533/−/− or Tg(OvPrP-V136)4166/−/− mice as sources of OvPrP-A136 and OvPrP-V136 substrates for protein misfolding amplification (PMCA) [25] using SSBP/1. While neither template spontaneously converted to PrPSc in the absence of seeded prions (Fig. 4A and E), SSBP/1 reproducibly converted OvPrP-V136 to OvPrPSc-V136 during a single round of PMCA (Fig. 4C). In contrast, conversion was not observed after a single round of PMCA when OvPrP-A136 was used as template (Fig. 4B and Fig. 5A). We therefore used serial PMCA (sPMCA) [26] over 10 rounds to address whether conversion of OvPrP-A136 to OvPrPSc-A136 might be detected after prolonged replication. Serial PMCA was performed in triplicate using equal amounts of PrP from three different Tg(OvPrP-A136)3533/+ or Tg(OvPrP-V136)4166/+ mice brains using sheep SSBP/1 as seed. Apart from a slight but consistent decrease between rounds two and five, OvPrPSc-V136 production, detected using mAb 6H4, was sustained throughout rounds one to 10 (Fig. 4C). As expected, mAb PRC5 failed to detect OvPrPSc-V136 (Fig. 4G). In contrast, mAb 6H4 detected OvPrPSc-V136 (Fig. 3B).

Figure 2. Analyses of PrPSc in the brains of SSBP/1 infected mice. In A and C, densitometric analysis of immunoblots was used to measure the amounts of protease-resistant OvPrPSc as a function of GdnHCl concentration. The dose-response curve was plotted using a Gaussian non-linear least-square fit. Each point represents the mean value derived from densitometric quantification of PK-resistant PrP in three diseased mouse brains. Error bars correspond to standard errors of the mean. A. Conformational stability analysis using mAb 6H4 of OvPrPSc-A136(S) in Tg(OvPrP-A136)3533/−/− mice (red line), OvPrPSc-V136(U) in Tg(OvPrP-V136)4166/−/− mice (blue line), and total OvPrPSc in Tg(OvPrP-A/V136) mice (black line). B. Black asterisks compare differences between OvPrPSc-A136(S) and total PrPSc in Tg(OvPrP-A136)3533/−/− and Tg(OvPrP-A/V) mice; blue asterisks compare differences between OvPrPSc-A136(S) in Tg(OvPrP-A136)3533/−/− mice and OvPrPSc-V136(U) in Tg(OvPrP-V136)4166/−/− mice. C. Conformational stability analysis using mAb PRC5 of OvPrPSc-A136(S) in Tg(OvPrP-A136)3533/−/− mice (solid line) and OvPrPSc-V136(U) in Tg(OvPrP-V136)4166/−/− mice (dashed line). **P<0.005, ***P<0.001. In B and D, representative immunoblots of PK-resistant PrP in the brains of three infected groups of Tg(OvPrP-A136)3533/−/−, Tg(OvPrP-V136)4166/−/−, and Tg(OvPrP-A/V136) mice using mAb 6H4 (B) and mAb PRC5 (C). Times of onset of disease for analyzed mice are also provided. OvPrP-V136 is indicated by blue symbols and text; OvPrP-A136 is indicated by red symbols and text. doi:10.1371/journal.ppat.1003692.g002

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OvPrP<sup>S<sub>c</sub>-A136</sup> was undetectable with mAbs 6H4 or PRC5 until round eight, after which levels decreased during rounds nine and 10 (Fig. 4B and F).

Forced templating of a dominant PrP<sup>S<sub>c</sub></sup> conformation in mice expressing both susceptibility alleles

Having established that Tg mice expressing OvPrPC-V136 and OvPrPC-A136 propagate SSBP/1-V136(U) and SSBP/1-A136(S) prions with relatively rapid and slow incubation times respectively, we produced Tg(OvPrP-A/V136) mice expressing both OvPrPC-A136 and OvPrPC-V136 and inoculated them with SSBP/1 to examine whether disease developed with fast, slow or intermediate kinetics. Although more rapid than the ~130 d onset of disease in Tg(OvPrP-A/V136) mice (P = 0.0094), the mean 105 ± 5 d onset of disease contrasted with the >400 d SSBP/1 incubation times observed in Tg(OvPrP-A136)3533<sup>+</sup>/2 mice. In accordance with this notion, histoblotting using mAb 6H4 showed that the neuroanatomical distribution of OvPrP<sup>S<sub>c</sub>-V136(U)</sup> in the brains of diseased Tg(OvPrP-A/V) mice mirrored the diffuse deposition of the OvPrP<sup>S<sub>c</sub>-V136(U)</sup> conformer located in similar sections of rapid incubation time Tg(OvPrP-V136)4166<sup>+</sup>/2 mice (Fig. 3C).

While the rapid SSBP/1 incubation times, and properties of the converted PrP<sup>S<sub>c</sub></sup> in diseased Tg(OvPrP-A/V) were consistent with propagation of SSBP/1-V136(U) prions, remarkably, western blotting of diseased Tg(OvPrP-A/V136) brain extracts with mAb PRC5 revealed substantial conversion of OvPrPC-A136 to OvPrP<sup>S<sub>c</sub>-A136</sup> (Fig. 2A), indicating that OvPrP<sup>S<sub>c</sub></sup> produced in Tg(OvPrP-A/V) mice shared the conformation of OvPrP<sup>S<sub>c</sub>-V136(U)</sup> produced in SSBP/1 infected Tg(OvPrP-V136)4166<sup>+</sup>/2 mice. In accordance with this notion, histoblotting using mAb 6H4 showed that the neuroanatomical distribution of OvPrP<sup>S<sub>c</sub>-A136</sup> in the brains of diseased Tg(OvPrP-A/V) mice contrasted with the diffuse deposition of the OvPrP<sup>S<sub>c</sub>-A136</sup> conformer located in similar sections of rapid incubation time Tg(OvPrP-V136)4166<sup>+</sup>/2 mice (Fig. 3C).

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OvPrP<sup>S<sub>c</sub>-A136</sup> was undetectable with mAbs 6H4 or PRC5 until round eight, after which levels decreased during rounds nine and 10 (Fig. 4B and F).

**Figure 3. Representative OvPrP<sup>S<sub>c</sub></sup> distribution in the CNS of diseased transgenic mice.** OvPrP-V136 and OvPrP-A136 are indicated by blue and red text respectively. Times of onset of disease (d) for individual mice analyzed in histoblots are provided. Sections through the midbrain, pons and oblongata are shown for SSBP/1 infected Tg(OvPrP-A136)3533<sup>+</sup>/2 mice (A and D); Tg(OvPrP-V136)4166<sup>+</sup>/2 mice (B and E); and Tg(OvPrP-A/V) mice (C and F). In panels G and H, sections were not treated with PK and show distribution of total PrP. Histoblots in panels A-G were probed with mAb 6H4; histoblots in D-H were probed with mAb PRC5.

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We then used mAb PRC5 to determine the conformation of OvPrPSc-A136 among total OvPrPSc produced in the brains of diseased Tg(OvPrP-A/V136) mice. The 1.64 GdnHCl1/2 value of OvPrPSc-A136 produced under these conditions was distinct from that of OvPrPSc-A136(S) produced in long incubation time Tg(OvPrP-A136)3533+/ mice with PK treatment; in C and G, the first lanes of each panel correspond to brain extracts from Tg(OvPrP-V136)4166+/- mice without PK treatment. In D and H, the first lanes of each panel correspond to mixtures of brain extracts from Tg(OvPrP-A136)3533+/ mice without PK treatment. In B and F, the first lanes of each panel correspond to brain extracts from Tg(OvPrP-V136)4166+/- mice without PK treatment. In B, C, and H, three different brains, indicated by #1–#3 were used as sources of substrates. In each case, the second lane of the second panel contains PK treated SSBP/1. In panels A and E, we probed histoblots of the CNS from diseased Tg(OvPrP-A/V136) mice with mAb PRC5 to assess the appearance and distribution of OvPrPSc-A136(U). In contrast to the punctate deposits of OvPrPSc-A136(S) in long incubation time Tg(OvPrP-A136)3533+/ mice (Fig. 3A and D), OvPrPSc-A136(U) in Tg(OvPrP-A/V136) mice (Fig. 3B) acquired a diffuse deposition and a distribution pattern that was equivalent to OvPrPSc-A136(U) in Tg(OvPrP-V136)4166+/- mice (Fig. 3B). Consistent with the co-expression of each allele in identical cell populations of Tg(OvPrP-A/V) mice, spatial distributions of 6H4- and PRC5-reactive PrP coincided in all analyzed sections of Tg(OvPrP-A/V) mice (Fig. 3G and H).

Figure 4. Detection of OvPrPSc by sPMCA. In A and E, the first two lanes of each panel correspond to brain extracts from Tg(OvPrP-A136)3533+/ and Tg(OvPrP-V136)4166+/- mice respectively without PK treatment. A136, V136, and A136/V136 refer to substrates from Tg(OvPrP-A136)3533+/ mice, Tg(OvPrP-V136)4166+/- mice, and mixtures of the two respectively. In B and F, the first lanes of each panel correspond to brain extracts from Tg(OvPrP-A136)3533+/ mice without PK treatment; in C and G, the first lanes of each panel correspond to brain extracts from Tg(OvPrP-V136)4166+/- mice without PK treatment; in D and H, the first lanes of each panel correspond to mixtures of brain extracts from Tg(OvPrP-A136)3533+/ and Tg(OvPrP-V136)4166+/- without PK treatment. In B-D and F-H, three different brains, indicated by #1–#3 were used as sources of substrates. In each case, the second lane of the second panel contains PK treated SSBP/1. Panels A and E are sPMCA using the indicated templates in the absence of a prion seed. Numbers in black above each lane refer to the round of sPMCA. C refers to a control in which no SSBP/1 seed was added. In each case C, and all numbered lanes were treated with PK. Blots were probed with mAbs 6H4 or PRC5 as indicated.

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PMCA reproduces strain specific effects of one allele product on another

To simulate the combined effects of OvPrP-A136 and OvPrP-V136 on PrP conversion in Tg(OvPrP-A/V136) mice in vitro, we mixed equal quantities of OvPrP-A136 and OvPrP-V136 in PMCA reactions seeded with SSBP/1. Under these conditions, similar to when OvPrP-V136 was present in isolation (Fig. 4C), we observed early, reproducible conversion to OvPrPSc in round one (Fig. 4D). Probing of western blots with mAb PRC5 showed that OvPrPSc-A136 was a component of this converted material (Fig. 4H). Thus, similar to our observations in Tg mice, the presence of OvPrP-V136 induced the relatively rapid conversion of OvPrP-A136 to OvPrPSc-A136 by SSBP/1. Interestingly, subsequent conversion of both OvPrP-A136 and OvPrP-V136 diminished in rounds two to five, ultimately becoming undetectable through rounds six to 10 (Figs. 4D and H).

SSBP/1 was originally produced from a pool of diseased sheep brains from the positive selection line in the Neuropathogenesis Unit (NPU) Cheviot sheep flock, and has subsequently been passed as a pool. We next compared the seeding properties of SSBP/1 with those of SSBP/1-A136(S) or SSBP/1-V136(U) prions derived from SSBP/1-infected Tg(OvPrP-A136)3533+/− or Tg(OvPrP-V136)4166+/− mice. We monitored conversion of OvPrP-A136 or OvPrP-V136 templates every two hours for a total of 12 h of PMCA. SSBP/1-V136(U) had the same PMCA properties as SSBP/1: both SSBP/1 and SSBP/1-V136(U) prions efficiently converted OvPrP-V136 in isolation, but not OvPrP-A136 in isolation; when both templates were present in the PMCA reaction, the presence of OvPrP-V136 facilitated conversion of OvPrP-A136 to OvPrPSc-A136 by SSBP/1 or SSBP/1-V136(U) prions (Figs. 5A and B). In contrast, SSBP/1-A136(S) prions converted either OvPrP-V136 or OvPrP-A136 templates to PrPSc when they were present in isolation, the latter being unequivocally confirmed to be OvPrPSc-A136 using mAb PRC5 (Fig. 5C); however, in the presence of both templates SSBP/1-V136(U) prion propagation was inhibited (Fig. 5C). The properties of prions derived from Tg(OvPrP-A/V) mice differed from SSBP/1, SSBP/1-A136(S) or SSBP/1-V136(U) prions. Like SSBP/1 and SSBP/1-V136(U), prions passaged through these mice efficiently converted OvPrP-A136 to OvPrPSc-A136, but not OvPrP-A136. However, unlike

![Figure 5. PMCA using defined seeds and substrates.](image_url)
SSBP/1 and SSBP/1-V136(U), such prions failed to facilitate conversion of OvPrPC-A136 in the presence of OvPrPC-V136 (Fig. 3D).

Discussion

Novel transgenic mouse models for analysis of the OvPrP-A/V136 dimorphism

Previous studies described the production of Tg mice expressing OvPrP, and reported their susceptibility to scrapie prions [27–32]. The most widely characterized models are tg338 mice expressing OvPrP-V136 [31], and Tgov59 [33] or Tgov4 [29] lines expressing OvPrP-A136. In the case of tg338 mice, the prion was comprised of a bacterial artificial chromosome insert of 125 kb of sheep DNA, while in the case of Tgov59 and Tgov4 mice the neuron specific enolase promoter was used to drive OvPrP expression. These lines are maintained on different heterogeneous genetic backgrounds, and CNS expression levels in tg338 mice are ~8–10-fold higher than wild type, while Tgov59 and Tgov4 lines each over express OvPrP-ARQ at levels ~2–to 4-fold higher than those found in sheep brain. Spontaneous neurological dysfunction has been reported in Tg lines over expressing OvPrP [27,31].

Tg(OvPrP-A136)3533+/− and Tg(OvPrP-V136)4166+/− mice express transgene-encoded PrP, either slightly lower, or slightly higher than PrP levels normally expressed in the CNS of wild type mice. Since Tg(OvPrP-A136)3533+/− and Tg(OvPrP-V136)4166+/− lines were produced using the cosSHa.Tet cosmid vector which drives expression from the PrP gene promoter [34], we expected expression of OvPrP–A136 and OvPrP–V136 in identical neuronal populations, and therefore that both alleles are co-expressed in the same cells of Tg(OvPrP-A/V) mice. Finally, other than variable transgene insertion loci, both lines are otherwise syngeneic on an inbred Pmng+/−/FVB background.

Previous studies reported on Tg mice expressing OvPrP with V at 136, referred to as Tg(OvPrP)14882+/− mice, that were also produced in a Pmng+/−/FVB background using the cosSHa.Tet cosmid vector [32]. However, in that study, comparable Tg mice expressing OvPrP-A136 were not reported. Median SSBP/1 scrapie incubation times in Tg(OvPrP-A136)3533+/− and Tg(OvPrP-V136)4166+/− mice were 75 d, and this line expresses OvPrP at levels only slightly higher than Tg(OvPrP-V136)4166+/− mice. While we exercise caution when comparing results from mice produced by different groups, the otherwise similar properties of Tg(OvPrP-A136)3533+/− and Tg(OvPrP-V136)4166+/− mice suggest that even slight differences in the levels of transgene expression can have significant effects on prion incubation time.

A clear link to codon 136 genotype and susceptibility/resistance to different sheep scrapie isolates has been described in multiple previous studies. Importantly, the influence of residue 136 on the transmission of SSBP/1 and CH1641 prions in Tg(OvPrP-A136)3533+/− and Tg(OvPrP-V136)4166+/− mice is in accordance with the properties of these isolates in sheep of various genotypes [17]. Generally, increased susceptibility to scrapie is associated with expression of OvPrP–V136, with A/A136 being the most resistant, and V/V136 the most susceptible genotypes. In the case of SSBP/1 incubation periods are ~170 days in V/V136 sheep, while transmission to A/A136 sheep is relatively inefficient, with no disease recorded after ~1000 days [33]. While SSBP/1 eventually transmits to Tg(OvPrP-A136)3533+/− mice with incubation times exceeding 400 days, the general effects of the A/V136 dimorphism on SSBP/1 transmission observed in sheep are recapitulated in Tg(OvPrP-A136)3533+/− and Tg(OvPrP-V136)4166+/− mice (Table 1). Similarly, CH1641, which propagates efficiently in A/A136 sheep [35], preferentially propagates in Tg(OvPrP-A136)3533+/− mice (Table 1). In previous studies, CH1641 transmitted to Tg(OvPrP-V136)4166+/− mice with an ~250 d mean incubation time [36].

Although SSBP/1 incubation times are prolonged in A/V136 compared to V/V 136 sheep [35], in our studies incubation times were shorter in Tg(OvPrP-A/V) than in Tg(OvPrP-V136)4166+/− mice. While the condition of A/V136 heterozygosity has not been previously modeled in Tg mice, this difference may result from double the levels of transgene expression in Tg(OvPrP-A/V) mice compared to Tg(OvPrP-A136)3533+/− and Tg(OvPrP-V136)4166+/− mice. Tg(OvPrP-A/V) mice were derived by mating Tg(OvPrP-A136)3533+/− with Tg(OvPrP-V136)4166+/− mice, and therefore express greater total levels of OvPrP than Tg(OvPrP-A136)3533+/− and Tg(OvPrP-V136)4166+/− mice (Fig. 1A). Since the levels of OvPrP–V136 are equivalent in Tg(OvPrP-V136)4166+/− and Tg(OvPrP-A/V) mice, and we show that OvPrP-A136 also becomes available for conversion, this situation results in more available substrate for conversion. While previous studies revealed an inverse correlation between transgene expression levels and prion incubation times in Tg mice [8], whether shorter incubation periods in Tg(OvPrP-A/V)136 mice than in Tg(OvPrP-V136)4166+/− mice reflect overall differences in PrP expression levels remains uncertain. Differences in scrapie pathogenesis between mice and sheep may also reflect the influence of additional factors on disease in the natural host including other PRNP polymorphisms [37,38], and different involvements of the lymphoreticular system in sheep compared to Tg mice.

Novel insights into the role of the A/V136 polymorphism on strain selection and prion propagation

Our observations in Tg mice expressing individual allele products suggested that rapid or prolonged SSBP/1 incubation times in Tg(OvPrP-V136)4166+/− and Tg(OvPrP-A136)3533+/− mice respectively, reflected preferential conversion by SSBP/1 prions of OvPrP–V136, rapidly producing a relatively unstable OvPrP–V136(U) conformation that was diffusely deposited in the CNS, compared to the slower conversion of OvPrP–A136 to the more stable OvPrP–A136(S) conformer which accumulated in the CNS with a punctate pattern (Figs. 1–3). Our results are consistent with the selection by the A/V136 dimorphism of SSBP/1 over SSBP/2 and SSBP/1-V136(U) prions in Tg(OvPrP-A136)3533+/− and Tg(OvPrP-V136)4166+/− mice respectively. We also show that PMCA recapitulates the influence of the A/V136 polymorphism on the kinetics of SSBP/1 propagation observed in Tg mice. The general conclusions from these studies agree with previously published assessments of the mechanism of conformational selection by distinct PrP primary structures [39,40].

Based on the rapid SSBP/1 incubation times in Tg(OvPrP-A/V136) mice, and shared conformational and distribution properties of OvPrP–A136 produced under these conditions with OvPrP–V136(U) in Tg(OvPrP-V136)4166+/− mice, we speculated that OvPrP–A136 played no part during the propagation of SSBP/1 prions in Tg(OvPrP-A/V) mice. To address this we used mAb PRC5 to exclusively monitor conversion of OvPrP–A136.
OvPrP<sup>Sc</sup>-V136(U) is formed by conversion of OvPrP<sup>F2</sup>-V136 by SSB/1 prions, the resulting unstable conformation induces rapid conversion of OvPrP<sup>F1</sup>-A136 to OvPrP<sup>Sc</sup>-A136(U). That this outcome is dependent on allele co-expression within the host is demonstrated by the inability of the OvPrP<sup>Sc</sup>-V136(U) conformer to template OvPrP<sup>Sc</sup>-A136 when it is expressed in isolation.

Effects of OvPrP genotype on the propagation of scrapie prions were not controlled during the isolation and propagation of SSB/1. Passage of SSB/1 through Tg mice therefore allowed us to generate prions composed solely of OvPrP<sup>F2</sup>-A136, OvPrP<sup>Sc</sup>-V136, or mixtures of both, and to draw additional conclusions about the effects of the A/V136 dimorphism of prion propagation using PMCA. Similar to our observations in Tg mice, SSB/1 failed to convert OvPrP<sup>F1</sup>-A136 to OvPrP<sup>Sc</sup>-A136 by PMCA, except in the presence of OvPrP<sup>F2</sup>-V136 (Figs. 4 and 5). These conversion properties are shared with SSB/1-A136(U) prions, but are distinct from SSB/1-A136(S) prions, which show facile conversion of both OvPrP<sup>F1</sup>-V136 and OvPrP<sup>F2</sup>-A136. These results suggest that the SSB/1-A136(U) is the dominant strain in the natural SSB/1 isolate. Multiple parameters could account for this, including, but not restricted to, the effects of OvPrP genotype, for example as a result of exclusive propagation in sheep of the V136/V136 genotype, route of infection in the infected sheep, and differential/selective prion replication in the lymphoreticular or central nervous systems of sheep. While our analyses indicate the presence of both OvPrP<sup>V1</sup>-V136 and OvPrP<sup>V2</sup>-A136 alleles in SSB/1 (Table 1), it is important to note that SSB/1 was derived from a pool of sheep brains of undefined genotypes. PCR approach precludes assessment of the extent to which alleles are present in a sample, raising the possibility that the one or other allele exists as a minor component in SSB/1.

Our findings also suggest that PrP<sup>Sc</sup> conformers may cross-inhibit PrP conversion. In case of SSB/1-A136(S)/prions, the presence of OvPrP<sup>F2</sup>-V136 inhibited PMCA of OvPrP<sup>F2</sup>-A136 (Fig. 5C). Also, while SSB/1 seeding of PMCA reactions containing mixtures of OvPrP<sup>F1</sup>-A136 and OvPrP<sup>F2</sup>-V136 resulted in robust, reproducible conversion to OvPrP<sup>Sc</sup>-A136 as early as round one (Fig. 4H), total PrP<sup>Sc</sup> production was ephemeral with subsequent PrP<sup>Sc</sup> formation diminishing during rounds two to five, and conversion ultimately becoming undetectable after round six. Since early PrP<sup>Sc</sup> conversion was sustained out to round 10 when OvPrP<sup>Sc</sup>-A136 was not produced (Fig. 4C and G), these results are consistent with inhibited conversion of OvPrP<sup>F1</sup>-V136 to OvPrP<sup>F2</sup>-V136 by OvPrP<sup>Sc</sup>-A136. While early (round one) PMCA conversion of PrP<sup>Sc</sup> by SSB/1 with either OvPrP<sup>F1</sup>-V136 or mixtures of OvPrP<sup>F1</sup>-V136 and OvPrP<sup>F2</sup>-A136 correlates with early onset of disease following SSB/1 infection of both Tg(OvPrP<sup>F1</sup>-V136) and Tg(OvPrP<sup>F1</sup>-A136/V136) mice, the subsequent inhibitory effects of OvPrP<sup>Sc</sup>-A136 observed in PMCA would be impossible to detect in vivo, since Tg(OvPrP<sup>F1</sup>-A136/V136) mice succumb to the lethal effects of early PrP<sup>Sc</sup> accumulation. Consistent with an inhibitory effect of OvPrP<sup>F1</sup>-A136(U) prions from Tg(OvPrP<sup>F1</sup>-A136/V136) mice, while they converted OvPrP<sup>F1</sup>-V136 in isolation, failed to convert OvPrP<sup>F1</sup>-A136 to PrP<sup>Sc</sup> in the presence of OvPrP<sup>F1</sup>-V136 (Fig. 3D). Thus, the properties of prions from this defined genetic background differ from SSB/1. We emphasize that, despite PCR data supporting the presence of OvPrP<sup>A136</sup> alleles in this isolate, SSB/1 was derived from sheep of undefined OvPrP<sup>V</sup> genotypes, rather than sheep with a defined heterozygous OvPrP<sup>F1</sup>-A136 genotype.

Wider implications for the mechanism of prion propagation

The inter-related effects of PrP primary and higher order structures on prion transmission were addressed in the Conformational Selection Model, which proposed that strains are composed of a range of PrP<sup>Sc</sup> conformers, or quasi-species, and that only a subset of PrP<sup>Sc</sup> conformations is compatible with each PrP primary structure [41]. While this model also took into account the effects of polymorphic variation on prion propagation, it did so only in the context of Tg mice expressing individual PrP<sup>A</sup> allele products. Transgenetic studies of the human codon 129 methionine (M)/valine (V) polymorphism, and the analogous codon 132 M/leucine (L) polymorphism in elk, indicated that these dimorphisms acted to restrict or promote the propagation of particular prion strains [39,40]. While the responses of Tg(OvPrP<sup>A136</sup>/3533<sup>S</sup>) and Tg(OvPrP<sup>V136</sup>/4166<sup>V</sup>/L) mice are consistent with this notion, that is selection of the U conformer by OvPrP<sup>V136</sup>, and the S conformer by OvPrP<sup>A136</sup>, our unprecedented ability to analyze allele specific conversion in infected Tg(OvPrP<sup>A136</sup>/3533<sup>S</sup>) mice reveals a more complex mechanism where mixtures of PrP variants may assist or inhibit the propagation of strains under various conditions. For example, SSB/1 or SSB/1-V136 prions that facilitate conversion of OvPrP<sup>F2</sup>-A136 to OvPrP<sup>F2</sup>-A136(U) only in the presence of OvPrP<sup>F2</sup>-V136. Expressed in isolation, conversion of OvPrP<sup>F2</sup>-A136 is favored by the OvPrP<sup>Sc</sup>(S) conformer. Our results demonstrate that co-expression of different polymorphic forms of PrP, which would be the norm in humans and animals, have profound effects on conformational selection of prion strains.

The results reported here address the molecular mechanisms associated with the phenomenon of prion strain over-dominance first observed by Dickinson and Outram [42], and subsequently reported in other settings involving co-expression of long and short incubation time PrP alleles [43]. While this phenomenon was reconciled at the time by the assumption that TSEs were caused by unidentified viral agents, our results now indicate that the suggestion raised by those studies, namely that over-dominance most likely resulted from physical interaction of allele products of the scrapie incubation time locus during infection, was prescient. Our results support a molecular mechanism involving cross-templating of an otherwise resistant allele product by a dominant prion conformer, in this case OvPrP<sup>Sc</sup>(U), which, we speculate, involves physical association of otherwise “susceptible” and “resistant” allele products. Consistent with the observations reported here, prion strain interference may also utilize similar mechanisms of conformational selection in a host expressing different PrP allele products infected with long and short incubation period strains with different PrP<sup>Sc</sup> conformational stabilities [44,45].

In conclusion, we have used a combination of transgenic, immunologic, and in vitro approaches to explore the mechanism by which PrP primary structure variations and the conformations enciphered by different prion strains interact to control TSE propagation. While our results support previous studies indicating that PrP susceptibility polymorphisms, expressed in isolation, act to restrict or promote the propagation of particular prion conformers, we now show that under conditions of allele co-expression a dominant conformer may alter the conversion potential of an otherwise resistant PrP polymorphic variant to an unfavorable prion strain. While such responses are analogous to the phenotypic expression of genetically determined heritable traits, dominant prion conformers act epigenetically by means of protein-mediated conformational templating. By expanding the range of possible conformations adoptable by a particular prion protein primary structure, such interactive effects provide a mechanism for promoting strain fitness, and, we speculate, strain diversification. While the precise number scrapie strains in sheep and goats remains uncertain, the description of at least 24 additional major sheep PRNP polymorphisms, and combinations
thereof, is likely to have a significant influence on strain diversity.

Materials and Methods

Ethics statement
All animal work was conducted according to the National Institutes of Health guidelines for housing and care of laboratory animals, and performed under protocols approved by the Colorado State University Institutional Animal Care and Use Committee, with approval number 11-2996A.

Transgenic mice

Sequences upstream of codon 44 of the OvPrP-A136 and V136 coding sequences were replaced with the corresponding sequence from mouse PrP. The resulting constructs contained the OvPrP coding sequence, except for addition of an extra residue for glycine at codon 31, and the mouse PrP N-terminal signal peptide instead of OvPrP signal peptide. Tg mice were generated by cloning the OvPrP-A136 and OvPrP-V136 expression constructs into the cosSHa.Tet cosmid vector [34], and microinjection of embryos from inbred Prnp$^{0/0}$/FVB mice. Tg founders were identified by PCR screening of genomic DNA isolated from tail snips. Founder mice were mated with inbred Prnp$^{0/0}$/FVB mice, and generally maintained with the transgene in the hemizygous state, with Tg mice identified by PCR screening of genomic DNA from weanlings. It was also possible to generate homozygous counterparts of each line, and Tg(OvPrP-A/V136) mice were generated by crossing homozygous Tg(OvPrP-V136)$^{+/+}$/Tg(OvPrP-A136)$^{+/+}$ mice with homozygous Tg(OvPrP-A136)$^{+/+}$/Tg(OvPrP-A136)$^{+/+}$ mice. We used immunoblotting and Western blotting with mAb 6H4 (Prionics, Schlieren, Switzerland) to estimate the levels of OvPrP expression. Tg mice subsequently shipped to and maintained in Edinburgh were crossed onto the Prnp$^{0/0}$/129Ola background [46].

Inocula

SSBP/1 originated as a homologue of three natural scrapie brains that were subsequently passaged mostly through Cheviot sheep at the Neuropathogenesis Unit (NPU), Edinburgh UK [15,16]. CH1641 is a naturally infected cheviot sheep from the NPU flock [17]. The presence of OvPrP-A136 or OvPrP-V136 alleles in these samples was ascertained by restriction fragment length polymorphism analysis of the PCR amplified PRNP coding sequences.

Ten % mouse brain homogenates (w/v) were prepared in PBS containing 150 mM NaCl, 1.0% Triton X-100, and the complete TM cocktail of protease inhibitors (Roche, Mannheim, Germany). Samples were clarified by brief, low-speed centrifugation. Protein concentrations of brain homogenates used as substrates for PMCA were adjusted to contain equivalent amounts of OvPrP-A136 or OvPrP-V136, based on the estimated relative levels of transgene expression. Substrates in which OvPrP-A136 and OvPrP-V136 were mixed were adjusted based on the estimated relative levels of transgene expression, so that approximately equal amounts of each allele product were present in the PMCA reaction. PMCA reactions were performed as described previously [20,47] at a seed to substrate ratio of 1:100. One cycle corresponded to 20 seconds of sonication followed by 30 minutes incubation at 37°C. Controls samples were incubated for the same duration at 37°C without sonication. Amplified and control samples were digested with PK at a final concentration of 0.33 μg/μl and analyzed on western blots using mAbs 6H4 or 5RC5.

Western blotting

Brain homogenates and cell lysates were digested with 100 μg/ml or 30 μg/ml of PK respectively (Roche, Mannheim, Germany) in cold lysis buffer for 1 h at 37°C. Digestion was terminated with phenylmethylsulfonyl fluoride at a final concentration of 2 μM. Samples were boiled for 10 min in the absence of B-mercaptoethanol [14] and proteins were resolved by SDS-PAGE and transferred to polyvinylidenedifluoride Immobilon (PVDF)-FL membranes (Millipore, Billerica, USA). Membranes were probed with primary mAbs followed by horseradish peroxidase–conjugated anti-mouse secondary antibody (GE Healthcare, Little Chalfont, UK). Protein was visualized by chemiluminescence using ECL Plus (GE Healthcare, Piscataway, USA) and an FLA-5000 scanner (Fujifilm Life Science, Woodbridge, USA).

Conformational stability assay

Brain homogenates containing 5 μg protein were incubated with various concentrations of guanidine hydrochloride (GuHCl) in 96-well plates for 1 h at room temperature. Samples were adjusted with PBS to a final concentration of GuHCl of 0.5 M and transferred onto nitrocellulose (Whatman GmbH, Dassel, Germany) using a dot blot apparatus. After two PBS washes, the membrane was air-dried for 1 h, then incubated with 5 μg/mL PK in 50 mM Tris-HCl, pH 8.0, 150 mM NaCl, 0.5% sodium deoxycholate, 0.5% Igepal CA-630 for 1 h at 37°C. PK was
inactivated with 2 mM PMSF. The membrane was incubated in 3 M guanidine thiocyanate in Tris-HCl, pH 7.5 for 10 min at room temperature. After four washes with PBS, the membrane was blocked with 5% nonfat milk in TBST for 1 h, and probed with mAbs 6H4 (1:20,000) or PRC5 (1:5,000) overnight at 4°C, followed by HRP-conjugated goat anti-mouse IgG secondary antibody. The membrane was developed with ECL Plus and scanned with GE image quant 4000. The signal was analyzed with ImageQuant TL 7.0 software.

Histoblotting
Histoblots were produced and analyzed according to previously described protocols [23]. Images were captured with a Nikon DMX 1200F digital camera in conjunction with Metamorph software (Molecular Devices).

References


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Author Contributions
Conceived and designed the experiments: GCT ES NH. Performed the experiments: ES JB KGB HEK SB SK NH. Analyzed the data: GCT ES JB. Wrote the paper: GCT ES JB.
44. Dickinson AG, Fraser H, McConnell I, Outram GW, Sales DI, et al. (1975)
Extraneural competition between different scrapie agents leading to loss of


129/Ola mice carrying a null mutation in PrP that abolishes mRNA production
are developmentally normal. Mol Neurobiol 8: 121–127.