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Both N-Terminal and C-Terminal Histidine Residues of the Prion Protein are Essential for Copper Coordination and Neuroprotective Self-Regulation.

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Abstract

The cellular prion protein (PrP^C) is comprised of two domains – a globular C-terminal domain and an unstructured N-terminal domain. Recently, copper has been observed to drive tertiary contact in PrP^C, inducing a neuroprotective *cis* interaction that structurally links the protein's two domains. The location of this interaction on the C-terminus overlaps with the sites of human pathogenic mutations and toxic antibody docking. Combined with recent evidence that the N-terminus is a toxic effector regulated by the C-terminus, there is an emerging consensus that this *cis* interaction serves a protective role, and that the disruption of this interaction by misfolded PrP oligomers may be a cause of toxicity in prion disease. We demonstrate here that two highly conserved histidines in the C-terminal domain of PrP^C are essential for the protein's *cis* interaction, which helps to protect against neurotoxicity carried out by its N-terminus. We show that simultaneous mutation of these histidines drastically weakens the *cis* interaction and enhances spontaneous cationic currents in cultured cells - the first C-terminal mutant to do so. Whereas previous studies suggested that Cu²⁺ coordination was localized solely to the protein's N-terminal domain, we find that both domains contribute equatorially coordinated histidine residue side chains, resulting in a novel

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Credit Author Statement

KMS, DAH and GLM conceived of the study. KMS, GLM wrote the manuscript. KMS, LT, BW, JTMK, NCU, MJP, RDB and GPR performed and analyzed the experiments. KMS, DAH and GLM edited the manuscript. GLM and DAH secured the funding.

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bridging interaction. We also find that extra N-terminal histidines in pathological familial mutations involving octarepeat expansions inhibit this interaction by sequestering copper from the C-terminus. Our findings further establish a structural basis for PrP^C's C-terminal regulation of its otherwise toxic N-terminus.

Keywords

PrP; PrP^C; A β 42; NMR; EPR

Introduction

Prion diseases, or Transmissible Spongiform Encephalopathies (TSEs), are a class of contagious, fatal neurodegenerative illnesses brought on by the misfolding of the endogenous prion protein (PrP)[1,2]. These diseases, which are part of a larger group of protein aggregation disorders including Alzheimer's and Parkinson's diseases, are an ongoing threat to human health, as well as to agricultural and wildlife animals.

Prion diseases are initiated by the misfolding of the prion protein (PrP) from its cellular form (PrP^C) into a beta sheet rich form (PrP^{Sc})[1,2]. Misfolded PrP^{Sc} aggregates deposit as fibrillar structures in brain tissue. The self-propagating nature of these aggregates has been extensively studied[2–4], but the mechanism of their neuronal toxicity is still unknown. PrP null mice do not exhibit symptoms of prion disease, suggesting that the disease is due to the gain of a toxic function, rather than the loss of essential activity[5]. PrP null mice injected with PrP^{Sc} do not show symptoms of prion disease, demonstrating the requirement of PrP^C as an essential precursor to PrP^{Sc}. In addition, PrP^C has been implicated as a signal transducer that transmits the toxicity initiated by PrP^{Sc}[6–9].

Several kinds of experiments suggest that alterations in PrP^C structure or conformation may mediate neurotoxic activities. For example, expression in transgenic mice of PrP molecules harboring deletions within the central domain of PrP, such as 105–125, causes a spontaneous neurodegenerative phenotype. In addition, monoclonal antibodies that bind the C-terminal domain of PrP^C are profoundly neurotoxic in vivo and in cultured cells[10–12]. It is thought that these manipulations eliminate the necessity for PrP aggregation, and directly induce PrP^C-mediated neurodegeneration. The effects of PrP deletions and anti-PrP antibodies can be measured in cell culture using patch-clamp recording, which detects the spontaneous, cationic currents that are induced[13].

In electrophysiological assays and in vivo, it has been observed that the flexible N-terminal domain, and in particular the first nine residues (23–31), are essential for neurotoxic activity[11,14]. Deletion of these residues, or treatment with antibodies and other ligands that bind this region, prevents the spontaneous currents and other neurotoxic activities[12,14]. In addition, cells expressing the N-terminal domain of PrP attached to another protein, EGFP, and entirely devoid of the C-terminal domain of PrP, exhibit spontaneous currents[11]. These results suggest that the N-terminal domain of PrP harbors a latent, toxic effector action, which is normally inhibited by interaction with the C-terminal domain.

The specific region on the C-terminus of PrP^C where toxic antibodies bind is near the sites of pathogenic, familial mutations[15,16]. Recently, our lab and others have shown that there is a *cis* interaction between the N-terminal and C-terminal domains of PrP^C, and that disruption of this interaction leads to neurotoxicity[15,17,18]. Taken together, these results suggest that an unregulated N-terminal domain of PrP^C is responsible for neurotoxicity, and that toxic PrP oligomers (ranging in size from ~5–10 PrP units[19,20]) may interfere with a natural protective regulation of the N-terminal domain by the C-terminal domain[11,18]. Despite the importance of PrP^C's self-regulation, only limited information is known about the essential regulatory interface between the two domains, although important insights have recently come from a study using a combination of cross-linking mass spectrometry and NMR[21].

In order to understand the molecular details of PrP^C's self-regulation, we consider the role of the protein's cognate metal ions. PrP physiology is highly dependent on the concentrations of divalent metal ions, particularly Cu²⁺. PrP has been shown to concentrate copper in brain tissue, and its expression is upregulated in response to increasing copper levels[22–24]. PrP is also known to regulate ion channels in a copper dependent manner[6,25]. Due to the strong link between copper and PrP function, the metal binding properties of the protein have been extensively studied[15,18,26–32]. In the N-terminus of the protein, there is an eight amino acid sequence repeated four times (octarepeat, OR; Figure 1). Each repeat contains one histidine, capable of coordinating the divalent metal ions Cu²⁺ and Zn²⁺. Cu²⁺ is found to bind the four histidine residues in the N-terminal domain's octarepeat region with sub-nanomolar affinity[29].

Recently, N-terminal copper binding has been observed to drive tertiary structure in PrP^C, inducing the aforementioned *cis* interaction between the N- and C-terminal domains of the protein[15,17,18,21]. The location of this interaction on the C-terminus overlaps with the sites of pathogenic mutations and toxic antibody docking. Copper delivered to cells has been shown to partially rescue toxicity resulting from deletion of a hydrophobic segment between the PrP^C N- and C-terminal domains[11]. Together, these results suggest that PrP^C's neuroprotective self-regulation is in fact driven by copper, which directly promotes the *cis* interaction between its toxic N-terminal and regulatory C-terminal domains.

How does copper cause the two domains to co-localize? Upon examination of the region on the C-terminus of PrP^C in which the copper-bound N-terminus docks, we were struck by the presence of two highly conserved histidine residues (Figure 1). The goal of this study is to determine if these C-terminal histidines facilitate the interdomain *cis* interaction, perhaps by acting as a tether to hold the two domains together. If so, a paradox is presented: if the copper ion is already coordinatively saturated by four N-terminal histidines, how could it bind additional C-terminal histidines?

To investigate a potential role of C-terminal histidines in the prion protein's *cis* interaction, we employed ¹H-¹⁵N HSQC NMR analysis of the murine protein. We show that mutation of histidines 139 and 176 (equivalent to human residues 140 and 177) results in a drastic weakening of the *cis* interaction. Furthermore, when His176 is deleted and a new histidine is added one turn up or down on the same helix, the location of the *cis* interaction tracks the

position of the histidine. We propose a model for this interaction in which copper acts as a tether – being simultaneously bound by both N and C-terminal histidine residues, analogous to the Cu(II) macro-chelate that occurs between His50 and Asp2 of α -Synuclein[33]. Using EPR spectroscopy, we find that the mechanism of this tether involves a histidine swap – one N-terminal histidine is removed from the copper ion and replaced with a C-terminal histidine. We show this model to be energetically plausible using molecular dynamics simulations and supporting density functional structure calculations. Together, these findings overturn the conventional view that Cu^{2+} is bound solely to the PrP^C N-terminal domain. Instead, both domains participate in metal ion coordination.

We find that, in addition to its structural effects, disruption of the *cis* interaction by mutation of both C-terminal histidines H139 and H176 enhances spontaneous ionic currents in cultured cells – the first C-terminal mutant to do this. Lastly, we find that the insertional mutations of extra, histidine containing octarepeat segments, which are linked to familial prion diseases, inhibit the protein's *cis* interaction by chelating copper away from the C-terminus. These findings advance our understanding of the toxicity caused by prion oligomers, toxic antibodies, and pathological mutations, and suggest that strengthening the *cis* interaction may be an effective therapeutic strategy for prion disease.

Results

Mutation of histidines 139 and 176 drastically weakens the prion protein's *cis* interaction

To examine whether C-terminal histidines contribute to the *cis* interaction, we mutated histidines 139, 176, and 186 of the murine prion protein to tyrosine, each in separate mutants. Because His186 is on the opposite face of the protein from the site of the *cis* interaction, we included it as a control. We chose tyrosine as a replacement amino acid because it is structurally similar to histidine, but cannot bind metals with similar affinity. At pH 7.4, the N-terminus of PrP^C binds copper in three coordination modes[26,27,29]. At low copper : PrP^C ratios, one copper ion is bound simultaneously by four histidines from the octarepeat region of PrP^C. At higher copper levels, multiple copper ions bind to the protein. In order to simplify interpretation of the data, we performed our experiments at pH 6.0, which locks the coordination into the first mode. Using ¹H-¹⁵N HSQC NMR at 37 °C and pH 6.0, we measured the peak intensities of C-terminal residues with no copper, and with one equivalent CuCl_2 . Dividing the intensity of each peak in the copper sample by its corresponding intensity in the apo sample, we calculated the intensity ratios (i/i_0) associated with the addition of one equivalence of copper to the protein, and repeated this process for each mutant. We then scaled the i/i_0 values to eliminate the effects of differential unspecific peak intensity reduction across mutants, and mapped the results onto the surface of the three-dimensional structure of the protein's helical C-terminus.

The wild type protein showed peak intensity reduction due in part to the paramagnetic relaxation effect (PRE), and also to exchange, in three areas as reported previously[17,18]. We refer to these areas of the helical C-terminal domain as “patch 1”, which is centered around His139, “patch 2”, which is centered around His176, and “patch 3”, which is located on helix three, between patch 1 and patch 2 (Figure 2). The peak intensity reduction in patches 1 and 2 is stronger than in patch 3, suggesting that they are closer to the bound

copper center of the OR. The H139Y mutation resulted in a much weaker peak intensity reduction at patch 1 than wild type, as did the H176Y mutation but at patch 2, while the H186Y mutation did not result in a difference to peak intensity reduction. The mutation of H186, which is the location of the pathogenic mutation H186R, appeared to affect C-terminal conformation of the protein and was therefore not further investigated. We varied protein and copper concentrations to test for the possibility of intermolecular interactions. We found that the peak linewidths were invariant as a function of protein concentration. In addition, the i/i_0 profiles were not affected as would be expected if two PrP^C molecules were forming a complex at higher concentrations. Together, these results suggest that there is only one PrP^C molecule involved per copper, and that the interaction is strictly between its N and C termini.

With the mutation of only one C-terminal histidine at a time, the *cis* interaction persisted, centered around the remaining histidine. We then made the double mutant (H139Y,H176Y), to test the theory that either histidine was capable of driving the interaction. In this mutant we observed that both patches 1 and 2 exhibited significantly decreased peak intensity reduction compared to wild type, abolishing the primary contacts of the established *cis* interaction (Figure 2). Peak intensity in patch 3 was also slightly *more* reduced than in wild type, suggesting a backup, albeit weak, mode of *cis* interaction. The strength of this backup interaction is best visualized using the bar plot of peak intensity ratios in Figure 2. We note that residues N-terminal to residue 120 are largely unstructured, resulting in peak overlap that confounds extraction of peak intensities, and have not been included in this analysis.

In order to be sure that these effects were due to the removal of histidines, and not due to addition of tyrosines which could sterically hinder the *cis* interaction, we performed these same experiments with the mutants (H139N,H176N) and (H139A,H176A), and observed the same results (data not shown).

We also performed this experiment on 90–230 PrP^C, which lacks the octarepeat histidines, to check that the observed C-terminal peak intensity reduction was due to a *cis* interaction rather than pure C-terminal copper binding. Although we observed slight intensity reduction around the C-terminal histidines in this mutant, the effect was much weaker than in the full-length protein. This observation provides convincing evidence that the reduction in peak intensity observed in the wild type protein upon the addition of copper results from the *cis*-interaction between its two domains, and not from transient copper binding solely to the C-terminal domain. Specifically, the loss of the octarepeat domain has a major effect on copper binding in the C-terminal domain, especially at His176.

Previous research by us and others has demonstrated the involvement of the non-octarepeat histidines H95 and H110 in copper binding[27,30,34]. At pH 7.4, copper coordination at these sites involves both the His side chain, as well as backbone amide nitrogens. To eliminate interference from coordination at these sites, we performed experiments at pH 6.0, which suppresses deprotonation of amides, thus reducing localized non-octarepeat copper binding. As a further control aimed at probing the involvement of histidines H95 and H110 in this interaction, we performed NMR on the construct (H95Y, H110Y). The strength and location of the *cis* interaction observed here is nearly identical to the wildtype protein

(Figure S1). We also performed continuous wave EPR on the wildtype protein at both pH 6.0 and pH 7.4. In the pH 7.4 spectrum we observed a previously characterized signal that arises from copper coordination from His95 and His110 due to backbone protonation, but we do not see this coordination in the pH 6.0 spectrum (Figure S2). Based on this data, we can rule out the possibility that the C-terminus is interacting with copper bound solely to a central region histidine complex. However, we cannot rule out the possibility that the C-terminus is interacting with copper bound to a complex formed of both the octarepeat and central region histidines.

Mutation of C-terminal residues in the region of the *cis* interaction surrounding histidines 139 and 176 does not abolish the interdomain interaction

In order to show that the *cis* interaction was driven by histidine-copper interactions rather than by a specific interaction between the two domains of the protein, we mutated most of the other residues in patches 1 and 2 spatially adjacent to the histidines to alanine. Specifically, the mutants (137A,138A,140A) and (172A,173A,175A,179A,180A) were created, as these are the residues that are located most closely to His139 and H176, and examined by NMR. The mutant (137A,138A,140A) adopted a misfold, preventing the data from being analyzed. The mutant (172A,173A,175A,179A,180A), which mutates the majority of patch 2 but leaves its histidine intact, maintained its i/i_0 profile consistent with the protein's *cis* interaction (Figure S3). These results reaffirm the importance of this C-terminal histidine in the copper driven *cis* interaction.

Deletion of histidine 176 and introduction of a new histidine one turn up or down its helix leads to a *cis* interaction localized to the new histidine

To show that the histidines were actively driving the *cis* interaction, rather than simply being positioned in the center of the involved areas, we deleted His176 and created two new mutants, one with a histidine one turn up and the other with a histidine one turn down its helix. As observed in Figure 3, the location of patch 2 follows the histidine. The ability of the histidine on helix 2 to control the location the *cis* interaction shows that it plays an active role in this docking event.

Electron paramagnetic resonance (EPR) experiments show no change in copper coordination environment, suggesting an equatorial histidine swap

Copper(II) in aqueous solutions exists in octahedral or square planer coordination geometries. With the knowledge that the four histidines of the PrP^C N-terminal octarepeat domain bind copper equatorially, this leaves only the axial binding sites available for a fifth, C-terminal histidine ligand. To investigate the coordination environment of the copper center, we employed X-band continuous wave (CW) EPR, Q-band electron nuclear double resonance (ENDOR), and hyperfine sub-level correlation (HYSCORE) spectroscopy at pH 6 (Figure 4).

As shown in Figure 4A, WT and the double mutant (H139Y, H176Y), which we refer to as "DM," produces the same X-band CW EPR spectra, which are well simulated by employing four equatorial nitrogen nuclei (^{15}N , $I = 1/2$) with a strong hyperfine coupling of a_{iso} ca. 55 MHz. This strong ^{15}N hyperfine coupling arises from directly coordinated nitrogens ($^{15}\text{N}_8$)

of the equatorial His imidazole rings. The remote histidine nitrogens ($^{15}\text{N}_e$), that are two bonds away from the $^{15}\text{N}_\delta$, typically have a much smaller hyperfine coupling. This smaller coupling was probed by using Mims-ENDOR shown in Figure 4B. Both WT and the double mutant (H139Y, H176Y) show only one set of weakly coupled ^{15}N , which is simulated using a hyperfine tensor A of [2.88, 2.88, 1.76] MHz, with $a_{\text{iso}} = 2.50$ MHz. This hyperfine tensor is determined by the simulation of both X-band and Q-band ^{15}N -HYSCORE spectra, as shown in Figure 4C and 4D (Note that both WT and DM show the same HYSCORE spectrum, see Figure 4D). The weak ^{15}N hyperfine interactions are assigned to the remote nitrogens $^{15}\text{N}_e$ of the equatorial imidazole rings,[35] as they are approximately 4.5% of that of the coordinated nitrogen, which is typical for imidazole-bound paramagnetic spin centers, such as imidazole-bound copper, vanadium and Rieske-type [2Fe-2S] clusters[35–37].

Besides the two sets of ^{15}N hyperfine interactions arising from the $^{15}\text{N}_\delta$ and $^{15}\text{N}_e$ of equatorially bound histidines, we did not observe other weak hyperfine interactions of ^{15}N as might be expected from a possible axial histidine. This suggests that there is no axial histidine ligand in WT protein. Indeed, we were able to show that the axial ligand is an exchangeable water ligand for both WT and the double mutant, as is evident from ^{17}O ($I = 5/2$) hyperfine interaction ca. 1.82 MHz observed from both ^{17}O HYSCORE (Figure 4D) and ENDOR (Figure 4E) experiment on the samples enriched with H_2^{17}O . This ^{17}O hyperfine interaction is consistent with that of an axial water ligand binding a Cu(II) center in the literature[38,39]. The axial water ligand is further confirmed by observing the exchangeable proton by Q-band Davies ENDOR (Figure 4F), which is simulated by using the ^1H hyperfine tensor A [-2.9, -2.8, 6.0] MHz, with the dipolar hyperfine T 2.95 MHz, corresponding to the distance of Cu-H ~ 2.99 Å.[40].

Taken together, these data show that there is no axial histidine binding in either the WT or the double mutant (H139Y, H176Y), and that copper coordination was unchanged by this mutation. This suggests that C-terminal histidines are in fact *not* acting as a fifth, axial ligand, but are instead displacing an equatorially bound N-terminal octarepeat histidine, giving equivalent coordination environments in both WT and DM proteins. We then produced the mutant (H84Y), which only has three octarepeat histidines. This mutant produced equivalent CW spectra to wild type, with four equatorial histidine ligands, and by NMR produced a slightly stronger *cis* interaction than in wild type, presumably because N-terminal histidines could no longer fully coordinate copper on their own (Figure S4). Combined with NMR evidence that a C-terminal histidine is driving the *cis* interaction, these data suggest that an N-terminal histidine is swapped out for a C-terminal histidine during the protein's *cis* interaction.

Pathogenic insertion and deletion mutation of histidine-containing octarepeat segments are explained by this model and supported by NMR experiments

An important class of pathogenic mutations in the prion protein involves the insertion of one or more extra copies of the histidine containing octarepeat segment. Mutations leading to extra octarepeats are associated with familial prion disease, and there is a drastic and sudden shift towards early onset disease when there are five or more extra octarepeat insertions[41]. To study these mutations, we employed the same NMR experiments listed above with one

equivalent of copper. As shown in Fig. 5, insertion of three extra histidine containing octarepeats did not inhibit the observed *cis* interaction, four extra histidines slightly weakened it, and five extra histidines almost completely inhibited the *cis* interaction. This suggests that insertion mutations sufficient to cause early onset prion disease may act by chelating copper away from C-terminal histidines and preventing the otherwise neuroprotective *cis* interaction. Alternatively, it is possible that the interaction is prevented by an increased entropic penalty due to the longer distance between the termini. In addition to insertional mutations, there are also deletion mutations in this region: deletion of one octarepeat segment does not cause prion disease, but deletion of two segments does[42–44]. This is consistent with the notion that three histidines from the octarepeat and one histidine from the C-terminus are involved in the *cis* interaction.

Mutation of C-terminal histidines enhances spontaneous currents

Certain mutations in the central region of PrP, such as 105–125 (referred to as CR) have been observed to cause spontaneous cationic currents in a variety of cell lines and in primary neurons[13]. The currents caused by CR mutations are suppressed by coexpression of wild type PrP, which mirrors the ability of coexpressed WT PrP to suppress the neurotoxicity of these mutations in transgenic mice[10]. C-terminal antibodies that are neurotoxic when injected into the mouse brain also cause spontaneous currents. We attribute the spontaneous currents and neurotoxicity observed in these scenarios to a weakening of the N-C inter-domain interactions within the PrP molecule[11,21]. We therefore tested whether the H139Y,H176Y mutations, by weakening the *cis* interaction, caused spontaneous currents in cultured cells.

Using whole-cell patch-clamping of N2a cells held at –90 mV, we observed spontaneous currents for (H139Y,H176Y), and not for a wild-type control (Figure 6). We used a –90 mV holding potential, which is more hyper-polarized than the normal resting potential of N2a cells (–70 mV), to enhance our ability to detect spontaneous currents; we have shown that these currents are voltage-dependent, being greater at more hyper-polarized membrane potential[11]. Consistent with studies of CR PrP mutants, the currents caused by H139Y,H176Y were suppressed by the addition of pentosan polysulfate (PPS), a polymer that sequesters the N-terminal domain of PrP[11].

Pyrenebutyrate (PB) is a polyaromatic, negatively charged small molecule that binds to positively charged groups such as the side chains of arginine and lysine. By shielding the positive charges, PB facilitates the ability of polybasic peptide regions, like those found at the N-terminus of PrP, to penetrate cell membranes[45–47]. We found (Figure 6b) that the presence of PB greatly enhanced the spontaneous currents caused the 105–125 mutation, even when recorded at –50 mV, a potential at which currents cannot normally be observed[11]. By using PB to enhance the detectability of spontaneous currents, we found that the (H139Y, H176Y) mutation caused currents at –70 mV, the normal resting of N2a cells, whereas no currents were detected at this potential for WT PrP (Figure 6b).

Protein molecular dynamics (MD) simulations and density functional structure calculations of copper binding in PrP^C

Histidines from both domains appear to be capable of cooperatively coordinating a copper ion, however, the specific His residues involved in this interaction and their location in the Cu²⁺ coordination sphere is not yet known. Starting from an initial coordination state where Cu²⁺ is bound by the four N-terminal His residues in a square planar coordination environment we identified eight additional possible configurations if one of the OR His residues is swapped for either His139 or His176. Using all-atom molecular dynamics simulations, we acquired a total of 675 ns of simulation time over all of the Cu-PrP^C configurations considered. We performed an additional seven simulations where two of the OR His residues are exchanged for His139 and His176 simultaneously (for a total of 285 ns of simulation time). While the specific atoms that comprise the Cu²⁺ coordination environment are different between each simulation, the number of atoms and the type of coordinating atoms immediately surrounding the Cu²⁺ center remain unchanged. We used the Cu²⁺ center and the imidazole heavy atoms (non-H-atoms) of the His residues that are involved in Cu coordination in each simulation along with the protein backbone heavy atoms of helix 2 and helix 3 as a common reference for accessing conformational stability. A clustering analysis was performed (details in methods section). Although there were no explicitly-bound water molecules at the Cu²⁺ axial positions, water molecules tended to occupy these sites for the majority of the simulation time, typically with no exchange or 1–2 waters exchanging during the course of the simulation. A persistent H-bond interaction was observed in several simulations between the interacting C-terminal Glu residues and one of these axially-retained waters. These results suggest that a His exchange between the N- and C-terminal domains holds the N-terminal OR region close to the C-terminal domain and that additional stabilization is afforded through electrostatic interactions between the cationic copper site and anionic residues in the C-terminus. Importantly, however, we also observe stable close-approach and these same stabilizing electrostatic interactions in the absence of His exchange, with the Cu²⁺ center bound by only the OR regions. This observation confirms that initial close approach of the Cu²⁺-bound OR region is already a favorable event and that His exchange then occurs as a further stabilizing consequence of the close association. An equilibrated, energy-minimized example of this interaction involving three OR His residues and C-terminal His176 simultaneously coordinating Cu²⁺ is shown in Figure 7.

To further characterize the His exchange event within the Cu²⁺ coordination environment we employed density functional structure calculations to assess the free energy differences between several potential exchange mechanisms. The His residues were modeled as 4-methylimidazole, representing the side chain, with the N_e-position protonated and the N_δ-position available for metal coordination. Axial waters were excluded from the [Cu²⁺ (4-methylimidazole)₄]²⁺ structure as these are only weakly bound and tend to form H-bonds instead of coordinate covalent interactions. We assessed two potential exchange mechanisms: i) an associative route, whereby the Cu²⁺ center coordinates a 5th imidazole moiety in an axial position, which we hypothesize would then displace an equatorial imidazole in a subsequent step, or ii) a dissociative route where one of the four imidazole moieties is first exchanged for a water molecule and then this water is subsequently

exchanged for another imidazole moiety (representing coordination of a C-terminal His residue). Comparing the relative free energies ($G_{(aq)}$) for the associative vs dissociative route, we find that both are relatively low, at +30.5 and +24.2 kJ/mol, respectively, suggesting that both mechanisms could contribute to ligand exchange. These estimates are from model coordination environments, however, and we anticipate that within the protein, the protein is likely to contribute significantly. While electrostatic interaction tends to draw the Cu^{2+} -bound N-terminal OR region toward the C-terminal domain, the steric bulk of the protein is likely to preclude formation of a crowded 5-His intermediate.

Discussion

Recent evidence shows that the flexible N-terminus of the prion protein acts as a toxic effector, and that this toxicity is down-regulated by interaction with the globular C-terminus[11,14]. Consistent with this model, deletions within the central region linking the two domains, as well as antibodies recognizing the region in the C-terminal domain that docks with the N-terminus, produce toxic effects in cultured neurons and in brain tissue. We hypothesize that these effects are caused by a disruption of the protein's self-regulation, which allows its N-terminus to extend away from the C-terminal domain, allowing aberrant interactions with the plasma membrane or membrane-bound receptors. NMR, EPR, and chemical cross-linking/MS studies strongly support a physical interaction between the N- and C-terminal domains of PrP[11,15,18,21].

Copper ions have been suggested to play a key role in mediating the interaction between the two domains of PrP, although the molecular details are not fully characterized. It is known that Cu^{2+} binds histidines in the octarepeat region of the protein's N-terminus, and that Cu^{2+} drives the *cis* interaction, but how does Cu^{2+} interact with individual amino acids to carry out this function? In this work, we show that two highly conserved histidines in the C-terminal domain of PrP are essential for the protein's *cis* interaction, and that their substitution with residues that do not bind copper diminishes the interaction, and also induces spontaneous ionic currents associated with an unregulated N-terminal domain. N- and C-terminal histidines work in synchrony to stabilize the *cis* interaction, suggesting that together they co-bind a copper ion, which serves as an inter-domain tether. Because the coordination environment of the copper is unchanged by this event, our data suggest that an N-terminal histidine is swapped with a C-terminal histidine, thereby stabilizing the *cis* interaction. An energy minimized example of this interaction is shown in Figure 7. Whereas previous PrP^C structural models suggested that Cu^{2+} coordination was limited to the protein's N-terminal domain, we find instead that both domains participate in binding the metal ion.

Similar, but rare, intra- and inter-domain copper-sharing mechanisms are observed in other protein systems. In blue copper proteins, copper links two domains, using three ligands from one domain and one ligand from another[48]. The copper chaperone Atx1 transfers copper to the Ccc2 ATPase through a transient intermediate in which the copper is co-bound by cysteines from both proteins[49]. A similar copper handoff occurs in the CusCBAF export pump in *E.coli*, and is mediated by methionine ligands[50]. Despite the existence of these interdomain and interprotein copper bridges in other protein systems, PrP^C has not

previously been known to display these structures. It is established that the N-terminus of PrP^C binds copper[26,27,29,30,51,52], and there is evidence in the literature that the C-terminus can as well[53–56]. However, this work is the first to demonstrate a non-local interaction, with a copper ion bridging N- and C-terminal histidine ligands and thereby stabilizing an inter-domain interaction. In this model, the affinity of Cu²⁺ for histidine residues in the N-terminal octarepeat region ($K_d=0.1$ nM), although high, is nevertheless weak enough to allow for exchange with histidine residues in the C-terminus.

The details of the *cis* interaction observed here provide a new explanation for a specific set of genetic prion diseases involving octarepeat variations. Individuals with at least five extra inserts of the octarepeat segment (nine total repeats) develop early onset prion disease. Based on the exceptionally high binding affinity between copper and this expanded OR segment ($K_d < 0.01$ nM), early onset disease was previously hypothesized to be due to either a loss of a copper dependent function of PrP^C, or a loss of copper mediated protection against misfolding[41]. In light of the new findings presented here, we suggest an alternative interpretation: the toxicity of these mutations may be due to the extra octarepeat histidines chelating copper away from C-terminal histidines, thereby disrupting the regulatory *cis* interaction. In addition, deletion of one octarepeat segment is nontoxic in humans, but deletion of two or more results in prion disease[42–44]. This is consistent with the notion that three histidines from the octarepeat and one histidine from the C-terminus are necessary for the *cis* interaction. Lastly, specific C-terminal point mutations that reduce negative charge and increase positive charge, such as D177N and E199K in mice (D178N and E200K in humans), are located close to H176, and therefore likely to reduce or prevent C-terminal copper binding, which would also impair the *cis* interaction.

The experiments described in this study have utilized a 1:1 Cu²⁺ : PrP^C stoichiometry. This stoichiometry is probably physiologically relevant, given the subnanomolar affinity of PrP^C for Cu²⁺ and the Cu²⁺ concentrations thought to be present at the synapse. However, we have shown previously that coordination of Cu²⁺ by PrP^C changes with concentration[29]. Studies into the nature of this interaction as a function of Cu²⁺ concentration may provide further insight into the *cis* interaction, PrP^C function and familial disease. Additionally, future investigations will consider zinc, another metal known to bind to PrP^C.

In summary, we have shown that two histidine residues in the C-terminal domain of PrP^C drive a protective *cis* interaction between the N- and C-terminal domains. We have designed a novel mutation, involving simultaneous deletion of these two histidines, that impairs N-C interaction and enhances spontaneous currents associated with neurotoxicity. Thus, our work furthers the body of evidence for a neuroprotective *cis* interaction in the protein. These findings suggest a novel PrP^C-mediated neurodegenerative mechanism whereby toxic prion oligomers, antibodies, and mutations elicit their effects through disruption of this interaction. Therefore, small molecule drugs or antibodies that stabilize and reinforce the *cis* interaction are potential therapeutic avenues to protect against toxicity in prion disease.

Materials and Methods

¹⁵N Labeled Protein Expression

Starting with the full length wild type mouse prion protein in the pJ414 vector backbone (DNA 2.0), we used QuickChange site-directed mutagenesis PCR to construct mutants, with primers generated manually and by www.primerdesigner.com. DNA was transformed into *E. coli* BL21Star (DE3) cells (Invitrogen) and expressed in M9 minimal media supplemented with 1 g/L ¹⁵N ammonium chloride (Cambridge Isotope Labs). Cells were grown at 37 °C until they reached an OD600 of 1.0, at which point the temperature was dropped to 30 C and protein expression induced with 1 mM isopropyl-1-thio-D-galactopyranoside (IPTG). Cells were allowed to express for 20 hours. Inclusion bodies were washed, then treated with 8 M guanidine HCl in pH 8 tris-acetate buffer to extract proteins. The extracted proteins were purified by immobilized metal affinity chromatography, allowed to fold overnight at pH 8, then transferred into pH 4.5 sodium acetate buffer by size exclusion chromatography. The proteins were then dialyzed into Milli-Q ultrapure water and purified further by reverse phase HPLC using C4 silica resin. Identity and purity were checked using a Sciex X500b mass spectrometer. The proteins were lyophilized, and quantified by UV-Vis spectrophotometry before use.

Nuclear Magnetic Resonance Spectroscopy

All samples were made to pH 6.0 in 10mM 2-(*N*-morpholino)ethanesulfonic acid (MES) buffer (Sigma), using potassium as a counterion and containing 10% D₂O. For all samples, the protein was added to a concentration of 300 μM. For samples with copper, CuCl₂ was used at 300 μM. ¹H-¹⁵N HSQC spectra were recorded at 37 °C on a Varian INOVA 600 MHz spectrometer (Varian, Santa Clara, CA) at UCSC NMR facility (Santa Cruz, CA), and on a Bruker 800 MHz spectrometer also at UCSC. NMR spectra were analyzed with NMR Pipe and Sparky using assignments transferred from previous experiments by visual inspection, and figures were made using Chimera and R. To determine a cutoff i/i_0 value to separate the residues involved in the *cis* interaction from the rest of the protein, we performed a kernel density estimation on the data using a Gaussian smoothing kernel. To eliminate the effects of differential unspecific peak intensity reduction across mutants, the data were scaled so that the center values of each mutant's group of unaffected peaks were aligned. We divided the residues into three categories based on their i/i_0 values: strongly affected (dark blue), weakly affected (light blue) and unaffected (grey). These divisions were created by using the local minimum separating the affected from unaffected residues in the wild type protein ($i/i_0 = 0.35$), and dividing the affected peaks into two groups ($i/i_0 = 0$ to 0.175, and $i/i_0 = 0.175$ to 0.35).

Electron Paramagnetic Resonance Spectroscopy

The X-band (9.38 GHz) continuous-wave (CW) EPR spectra were recorded on a Bruker (Billerica, MA) EleXsys E500 spectrometer equipped with a super-high Q resonator (ER4122SHQE). Cryogenic temperatures were achieved and controlled using an ESR900 liquid helium cryostat in conjunction with a temperature controller (Oxford Instruments ITC503) and gas flow controller. CW EPR data were collected under slow-passage, non-saturating conditions. The spectrometer settings were as follows: temperature = 15 K,

conversion time = 40 ms, modulation amplitude = 0.8 mT, and modulation frequency = 100 kHz; other settings are given in corresponding figure captions. Simulations of the CW spectra and the following pulsed EPR spectra were performed using EasySpin 5.1.10 toolbox[57,58] within the Matlab 2014a software suite (The Mathworks Inc., Natick, MA).

Q-band (~34.0 GHz) pulse ENDOR experiments were performed on a Bruker Biospin EleXsys 580 spectrometer equipped with a 10 W amplifier and a R.A. Isaacson cylindrical TE₀₁₁ resonator in an Oxford CF935 cryostat. ENDOR measurements were performed at 20 K by employing the Mims pulse sequence ($\pi/2$ - τ - $\pi/2$ -RF- $\pi/2$ - τ -echo) for small hyperfine couplings[59] or Davies pulse sequence (π -RF- $\pi/2$ - τ - π - τ -echo) for larger hyperfine couplings[60]. ENDOR spectra were collected stochastically by randomly hopping the RF excitation frequency[61]. Pulse sequences were programmed with the PulseSPEL programmer via the Xepr interface.

For Mims-ENDOR experiments,[59] the ENDOR intensities are modulated by the response factor (R) which is a function of the hyperfine coupling A and the time interval (τ) between the first and the second $\pi/2$ microwave pulse in the three-pulse sequence: $R \sim [1 - \cos(2\pi A\tau)]$. When $A\tau = n$ ($n = 0, 1, 2, 3 \dots$), this factor will be zero, corresponding to a minima in the ENDOR response, i.e., the hyperfine “suppression holes” in Mims-ENDOR spectra. This Mims-hole effect can be avoided by adjusting the τ value, as shown in Figure 4B.

X-band and Q-band HYSORE spectra were recorded on the Bruker Biospin EleXsys 580 spectrometer with a split-ring (MS5) resonator at 20 K using the pulse sequence $\pi/2$ - τ - $\pi/2$ - t_1 - π - t_2 - $\pi/2$ - τ -echo. The pulse length for inversion pulse (t_π) and the $\pi/2$ pulse ($t_{\pi/2}$) was 32 ns and 16 ns, respectively. Eight-step phase cycling was used. Time-domain spectra were baseline-corrected (third-order polynomial), apodized with a hamming window, zero-filled to eight-fold points, and fast Fourier-transformed to yield the frequency-domain spectra.

Whole-Cell Patch Clamp Experiments

Recordings were made from N2a cells 24–48 hr. after transfection. Transfected cells were recognized by green fluorescence resulting from co-transfection with pEGFP-N1. Whole-cell patch clamp recordings were collected using standard techniques. Pipettes were pulled from borosilicate glass and polished to an open resistance of 2–5 megaohms. Experiments were conducted at room temperature with the following solutions: internal, 140 mM Cs-glucuronate, 5 mM CsCl, 4 mM MgATP, 1 mM Na₂GTP, 10 mM EGTA, and 10 mM HEPES (pH 7.4 with CsOH); external, 150 mM NaCl, 4 mM KCl, 2 mM CaCl₂, 2 mM MgCl₂, 10 mM glucose, and 10 mM HEPES (pH 7.4 with NaOH). Current signals were collected from a Multiclamp 700B amplifier (Molecular Devices, Sunnyvale, CA), digitized with a Digidata 1440 interface (Molecular Devices), and saved to disc for analysis with PClamp 10 software.

Molecular dynamics simulations

The model of the Cu²⁺-bound mouse PrP^C protein was constructed from a representative structure from the NMR-derived dataset for 1XYX[62] (mouse PrP^C fragment 121–231). Residues 23–120 were added, with the ϕ and ψ angles for residues 57–91, the OR region,

two dominant and closely-related conformations that represented 72% of all structures sampled. Close inspection of the dominant clusters revealed that most simulations formed stabilizing electrostatic interactions between the Cu²⁺ coordination site and Glu207 and Glu211 in the C-terminal domain.

Density functional structure calculations

Density functional calculations employed Gaussian 16, revision A.03[67], with the B3LYP functional method[68] employing the 6–311+G(d,p) basis set on all atoms, and included the Grimme-D3 dispersion correction[69] for geometry optimizations and subsequent harmonic frequency calculations. The long-range stabilizing influence of solvation was calculated from the gas-phase optimized geometries using a self-consistent reaction field dielectric continuum (IEFPCM)[70] with $\epsilon = 78.36$ (for water) and employing the united atom topological model radii (UAHF). Relative free energies between calculated structures followed previously-published methods[64].

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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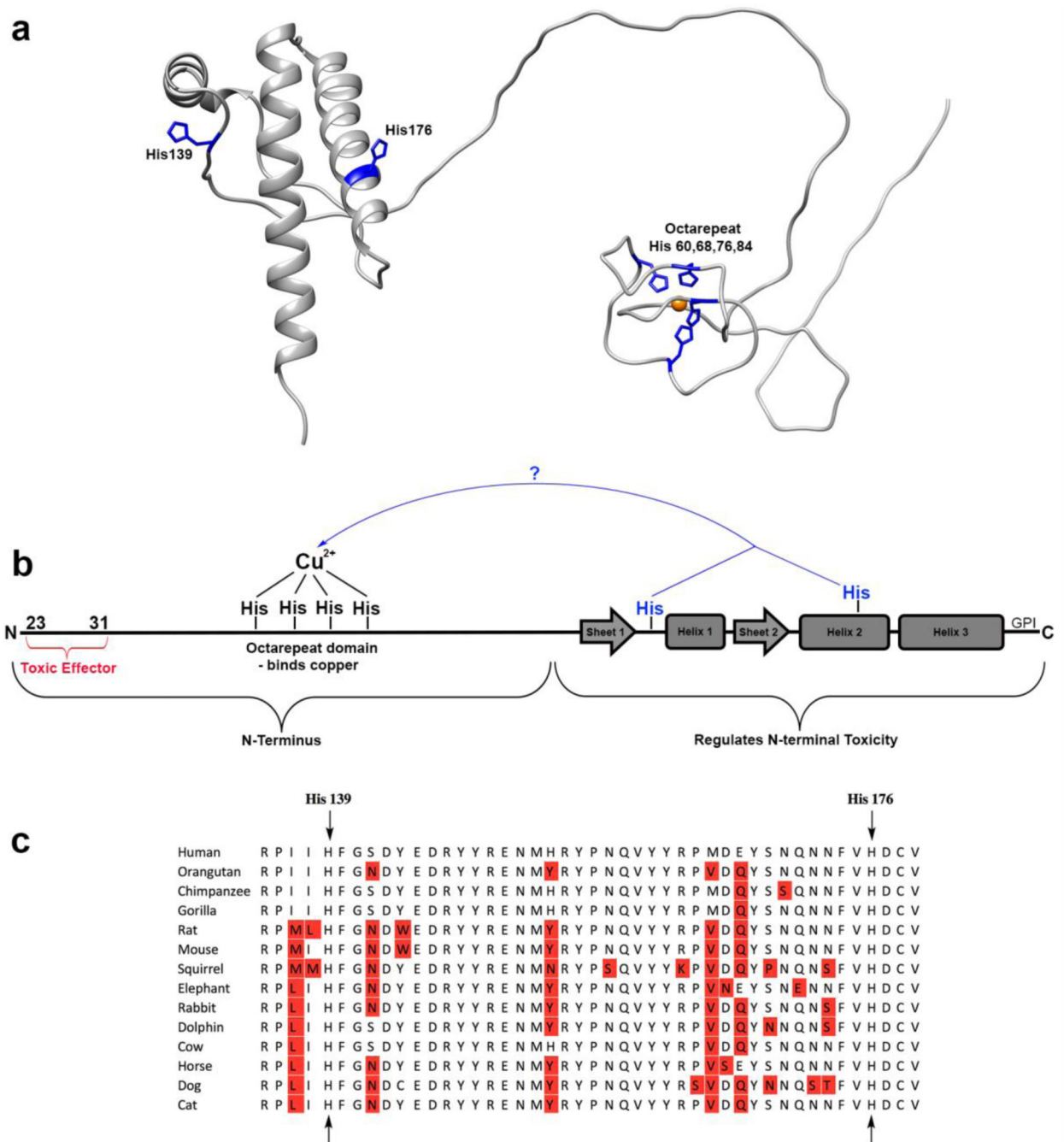
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Highlights

- The two domains of the cellular prion protein are tethered together by a copper ion
- This tethering is the basis of the protein's neuroprotective *cis* interaction
- Disruption of this interaction explains the toxicity of prion diseases
- Disruption may explain prion protein mediated toxicity in Alzheimer's disease

**Figure 1:**

a) Our current understanding of the prion protein. b) The potential new mechanism of self-regulation investigated in this paper: C-terminal histidines coordinating an N-terminally bound copper ion. c) Conservation of the histidines hypothesized to be involved. Residues with sequence variation are noted in red.

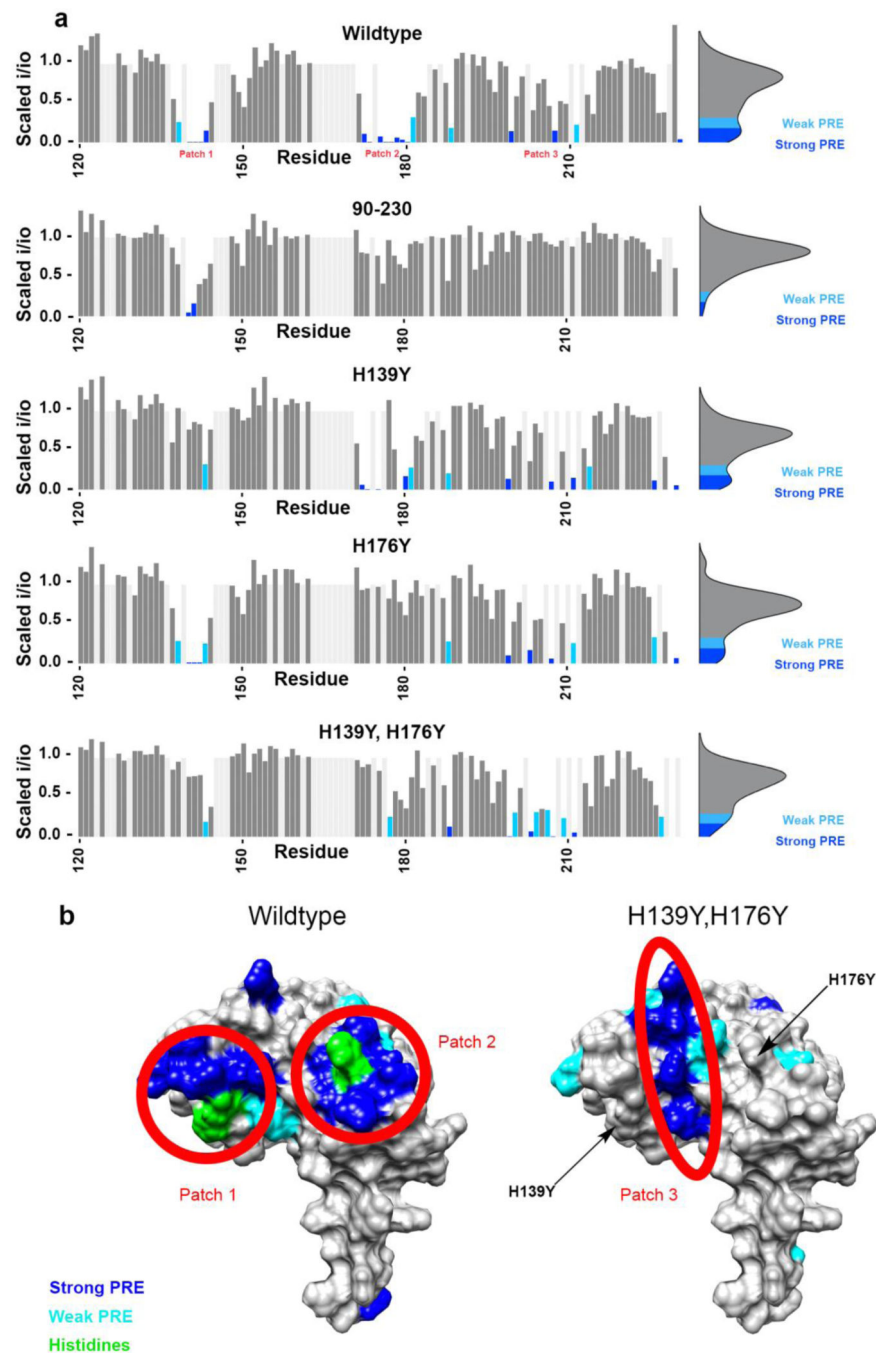


Figure 2.
a) Bar plots showing the magnitude of the peak intensity reduction derived from the paramagnetic relaxation effect (PRE) on specific residues of the protein's C-terminal domain, with 1 equivalent of CuCl_2 at pH 6.0 in 10 mM MES buffer for all five protein samples. Patches 1 and 2, which are located around histidines 139 and 176 respectively, are much more strongly affected in the wild type protein than in (H139Y,H176Y) and in 90–230 PrP^C. Unassigned residues are light grey and are drawn with no intensity reduction. **b)**

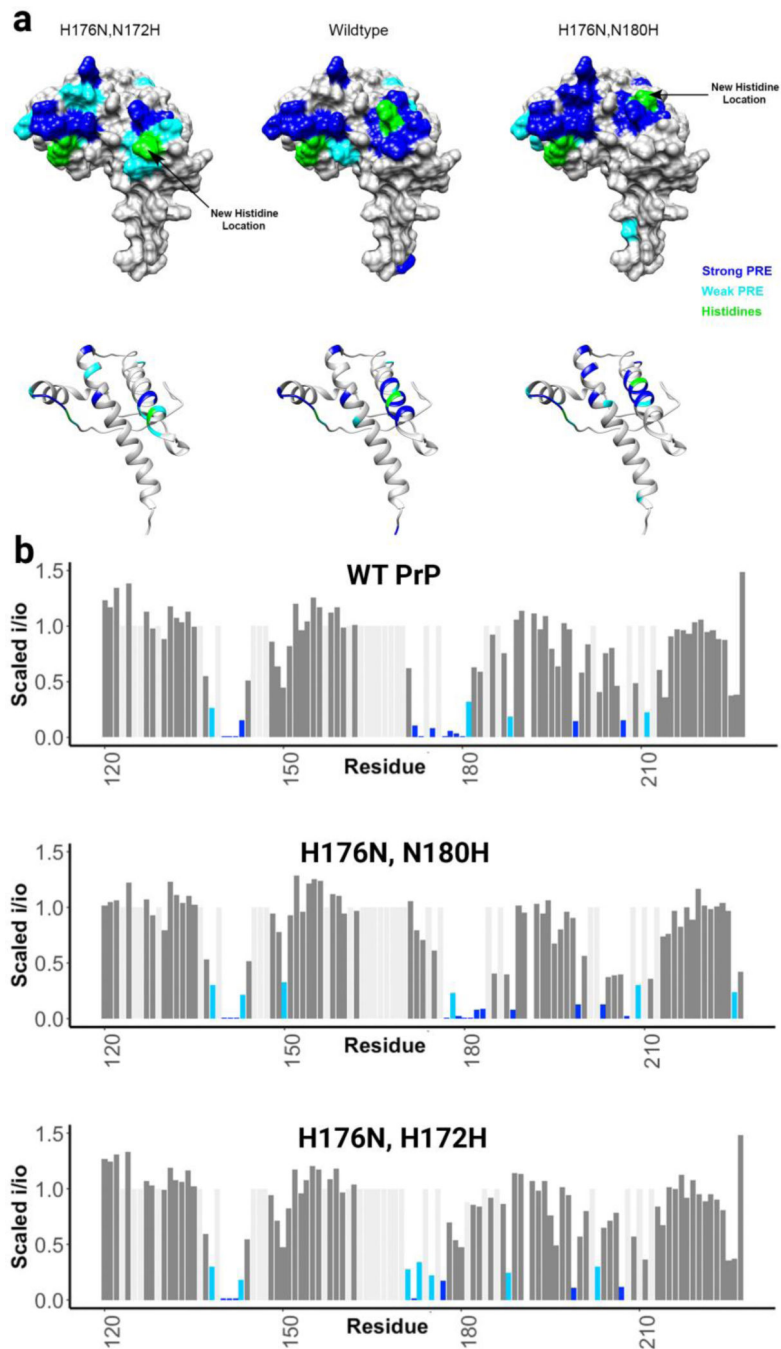
Surface representations showing areas engaged in the *cis* interaction, as measured by the intensity reduction of NMR cross peaks, plotted on PDB: 1XYX.

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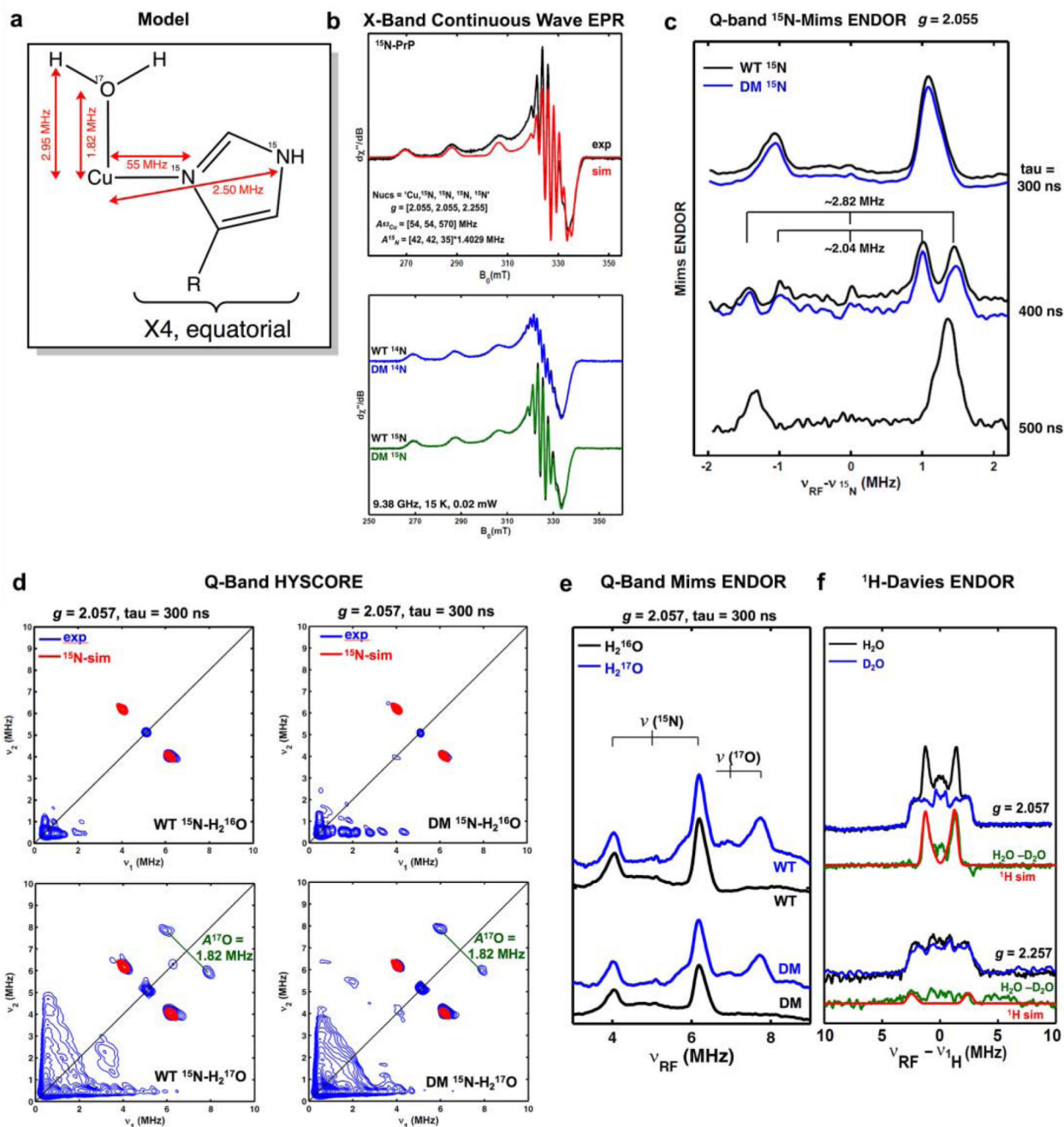
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**Figure 3:**

a) Surface representations showing areas engaged in the *cis* interaction, as measured by the intensity reduction of NMR cross peaks, plotted on PDB: 1XYX. This figure shows that the *cis* interaction follows histidine **b)** Bar plots showing the magnitude of the peak intensity reduction derived from the paramagnetic relaxation effect (PRE) on specific residues of the protein's C-terminal domain, with 1 equivalent of CuCl_2 at pH 6.0 in 10 mM MES buffer for all protein samples.

**Figure 4:**

a) Representation of the copper site. b) X-band continuous wave EPR spectra of WT PrP^C in agreement with simulation based on four equatorial nitrogen ligands and comparison of WT PrP^C to (H139Y, H176Y) PrP^C (labelled DM, for double mutant). c) Q-band ¹⁵N-Mims ENDOR showing weakly hyperfine interaction for both WT and DM. Varying tau values were used for confirming the Mims-hole effect. d) Q-band HYSCORE EPR spectra showing weakly coupled remote ¹⁵N from histidine and axial ¹⁷O from water. e) Q-band ¹⁷O Mims

ENDOR, confirming axial water coordination. f) ^1H -Davies ENDOR showing the disappearance of an exchangeable proton from water in when D_2O is used as solvent.

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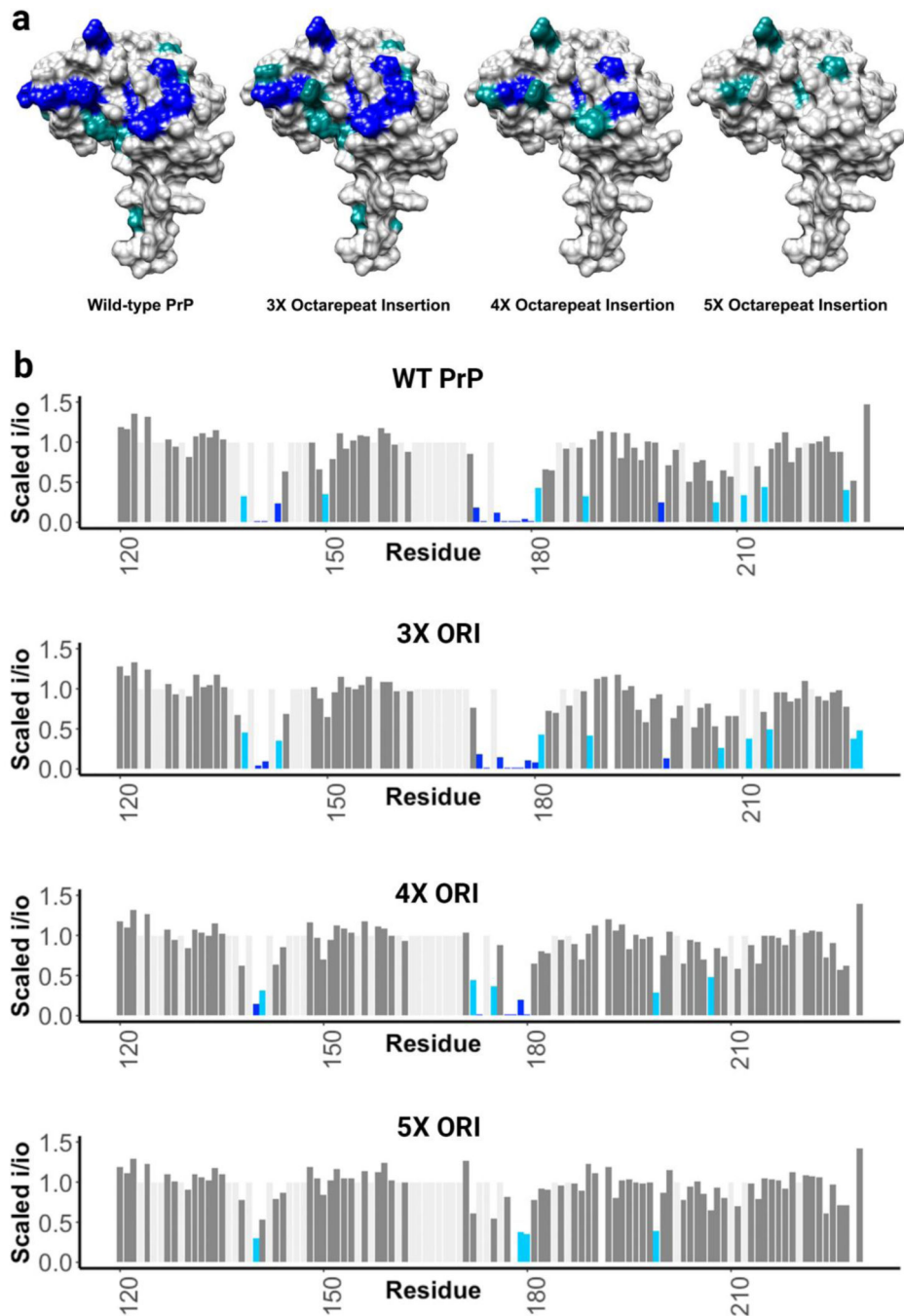


Figure 5.

a) Surface representations showing areas engaged in the *cis* interaction, as measured by the intensity reduction of NMR cross peaks, plotted on PDB: 1XYX. Five additional octarepeat regions results in a strong decrease in the *cis* interaction. **b)** Bar plots showing the magnitude of the peak intensity reduction derived from the paramagnetic relaxation effect (PRE) on specific residues of the protein's C-terminal domain, with 1 equivalent of CuCl_2 at pH 6.0 in 10 mM MES buffer for all protein samples.

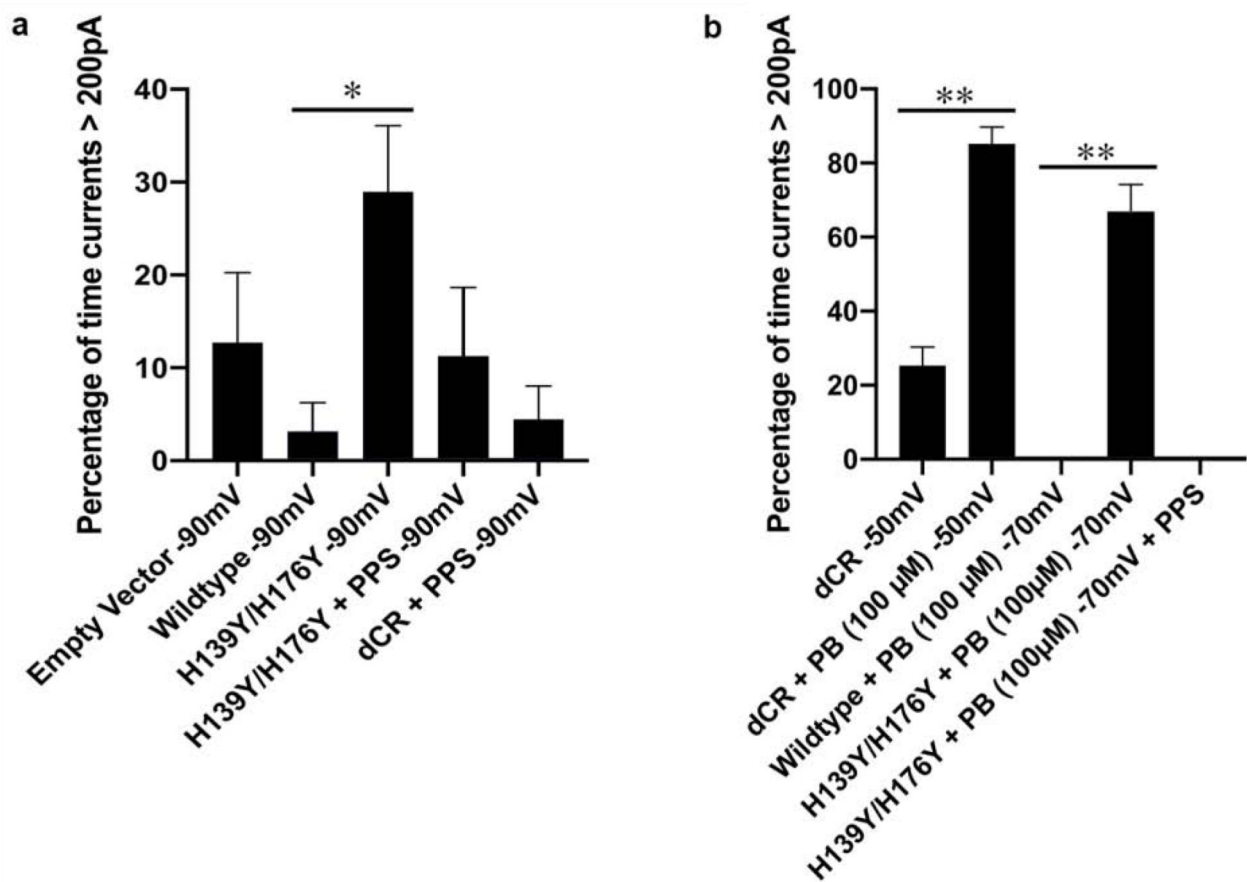


Figure 6.

a) Comparison of cells transfected with empty vector (N=10), wild-type PrP (N=5), (H139Y,H176Y) PrP (N=20), and CR PrP (N=10) showing that the H139Y,H176Y mutations significantly enhance spontaneous currents at a hyperpolarized holding potential (-90 mV) (* $p=0.045$, $\alpha=0.05$, one tailed two sample t-test), and these currents are suppressed by the addition of pentosan polysulfate (PPS). **b)** Comparison of cells expressing wild-type PrP (N=10), (H139Y,H176Y) PrP (N=10), and CR PrP (N=9) showing that the H139Y,H176Y mutations significantly enhance spontaneous currents at -70 mV in the presence of pyrenebutyrate (** $p<0.00001$), and these currents are suppressed by pentosan polysulfate. The CR mutation enhances currents in the presence of pyrenebutyrate, even at -50 mV (** $p<0.00001$). Error bars show the standard error of the mean.

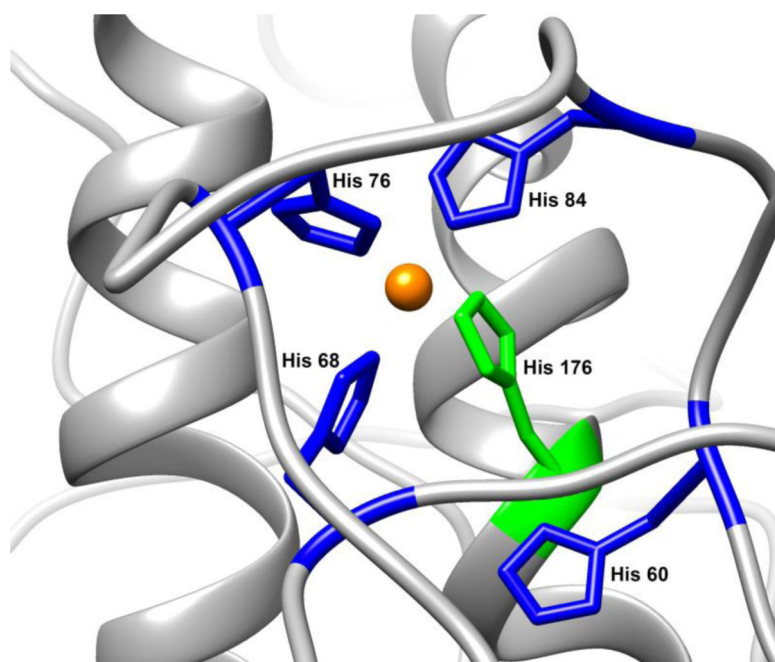


Figure 7:
A structural explanation for the results of this paper. An N-terminal histidine (blue) swaps out for C-terminal His176 histidine (green) during the protein's *cis* interaction, creating a copper tether between the two domains.

Table 1:

Summary of couplings from EPR experiments and their meanings.

Hyperfine Coupling	Origin	Conclusion
a_{iso} : 55 MHz in CW	Coordinated nitrogen on equatorial histidine	
a_{iso} : 2.50 MHz in Mims ENDOR and HYSORE	Remote nitrogen on equatorial histidine	-
T : 2.95 MHz in Davies ENDOR	Exchangeable proton of axial water	-
A : 1.82 MHz in HYSORE and Mims ENDOR	Oxygen from H_2^{17}O	Presence of axial water
Absence of weak coupling of ^{15}N in Mims ENDOR	Axial histidine ligand	No axial histidine

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Table 2:

Constructs tested by NMR in this paper

Construct	<i>Cis</i> Interaction
Wild type Mouse PrP ^C	<i>Cis</i> Interaction, four his ligands
H139Y	Interaction shifted towards H176
H176Y	Interaction shifted towards H139
H186Y	Interaction maintained
H186R	Interaction maintained
H139Y,H176Y	Strongly reduced interaction
H139N,H176N	Strongly reduced interaction
H139A,H176A	Strongly reduced interaction
M139A,I138A,F140A	Misfold / no data on interaction
N172A,N173A,V175A,V179A,N180A	Interaction maintained
Segment 90–230	No interaction
H176N,N172H	Interaction moved down helix
H176N,N180H	Interaction moved up helix
3XORI	Very weakly reduced interaction
4XORI	Weakly reduced interaction
5XORI	Strongly reduced interaction
H84Y	Slightly stronger <i>cis</i> interaction than WT
H95Y, H110Y	No difference in <i>cis</i> interaction from WT