Copper Alters Aggregation Behavior of Prion Protein and Induces Novel Interactions between Its N- and C-terminal Regions

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Background: Role of copper as an attenuator or facilitator in prion diseases is controversial.

Results: Copper-bound PrP does not aggregate at physiological temperature and shows two novel interactions between its N- and C-terminal domains.

Conclusion: Copper may act as an attenuator in prion diseases and induces novel long range inter-domain interactions in PrP.

Significance: This study might help in understanding the role of copper in prion diseases.

Copper is reported to promote and prevent aggregation of prion protein. Conformational and functional consequences of Cu2+-binding to prion protein (PrP) are not well understood largely because most of the Cu2+-binding studies have been performed on fragments and truncated variants of the prion protein. In this context, we set out to investigate the conformational consequences of Cu2+-binding to full-length prion protein (PrP) by isothermal calorimetry, NMR, and small angle x-ray scattering. In this study, we report altered aggregation behavior of full-length PrP upon binding to Cu2+. At physiological temperature, Cu2+ did not promote aggregation suggesting that Cu2+ may not play a role in the aggregation of PrP at physiological temperature (37°C). However, Cu2+-bound PrP aggregated at lower temperatures. This temperature-dependent process is reversible. Our results show two novel intra-protein interactions upon Cu2+-binding. The N-terminal region (residues 90–120 that contain the site His-96/His-111) becomes proximal to helix-1 (residues 144–147) and its nearby loop region (residues 139–143), which may be important in preventing amyloid fibril formation in the presence of Cu2+. In addition, we observed another novel interaction between the N-terminal region comprising the octapeptide repeats (residues 60–91) and helix-2 (residues 174–185) of PrP. Small angle x-ray scattering studies of full-length PrP show significant compactness upon Cu2+-binding. Our results demonstrate novel long range inter-domain interactions of the N- and C-terminal regions of PrP upon Cu2+-binding, which might have physiological significance.

Prion protein (PrP) is involved in transmissible spongiform encephalopathies, a group of diseases such as Creutzfeldt-Jakob disease, Kuru, Fatal Familial Insomnia, and Gerstmann-Strassler-Scheinker syndrome, characterized by neurodegeneration, spongiform cerebral tissue, and amyloid plaques. Such diseases are caused by the conformational transition of α-helix-rich cellular prion protein (PrPc) to the β-sheet-rich scrapie prion protein (PrPSc) (1). Copper ions (Cu2+) appear to have a significant role in such conformational transition (2). PrP exhibits strong binding toward Cu2+ with nano- to femtomolar affinity (3–5). Cu2+-binding sites are reported in the flexible N-terminal region (residues 23–120); four of them are present in the octapeptide repeats (residues 60–91) and one located at His-116 and His-111 (6–8) (residues are numbered according to the human PrP sequence). An additional Cu2+-binding site at His-187 has also been recognized in the C-terminal region of PrP (9–12).

Although the functional implication of Cu2+-binding to PrP is not yet clear, it is believed that Cu2+ is an important factor (as a facilitator or an attenuator) in prion diseases. Cu2+-binding induces conformational changes leading to reduction in the helical content and an increase in the β-sheet content especially in the region 90–120 (13), which partially overlaps with the well established amyloidogenic stretch 106–126 (14). Cu2+ has also been shown to induce conformational changes and increased stability in the octapeptide repeats of PrP (15, 16).

Cu2+ is shown to convert PrPSc from brain homogenates (2) and aged recombinant PrP (17) to protease-resistant and detergent-insoluble aggregates. However, these aggregates are structurally distinct from the scrapie prion protein (PrPSc). Expansion of octapeptide repeats has been suggested to cause an early onset of the prion disease (18, 19). In contrast, several studies show compelling evidence on the inhibitory role of Cu2+ on the amyloid formation of PrP in vitro (20–23). In vivo studies of Cu2+-binding to PrP are also not unequivocal. One study

The on-line version of this article (available at http://www.jbc.org) contains supplemental Figs. S1–S6 and Table S1.

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3 The abbreviations used are: PrP, prion protein; SAXS, small angle x-ray scattering; ITC, isothermal calorimetry; EOM, ensemble optimization method.
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reports the delay in the onset of prion disease upon removal of Cu\(^2+\) by chelation (24), whereas a different study shows that Cu\(^2+\)-depleted diet enhances prion disease (25). Thus, the nature and consequences of interaction of Cu\(^2+\) with PrP, both in vitro and in vivo, appear to be a complex phenomenon that remains to be investigated.

Conformational studies on binding of Cu\(^2+\) to PrP in vitro have been performed using peptides (26–28) or truncated variants of PrP (29–31) and at nonphysiological temperatures (2, 10, 13–17) and in water or buffers of low ionic strength (3, 9, 13, 32–34). We have used full-length PrP(23–231) to investigate conformational changes and aggregation upon Cu\(^2+\)-binding at physiological temperature (37 °C). We find that Cu\(^2+\)-bound PrP undergoes unusual temperature-dependent reversible aggregation. Interestingly, it aggregates at lower temperatures and resolubilizes as the temperature is raised. Our investigations using isothermal calorimetry (ITC), NMR, and small angle x-ray scattering (SAXS) provide insight into thermodynamic and conformational aspects of this process. Based on our study, we propose the involvement of the interaction of helix-2 and the octapeptide repeats region in the aggregation process. Our results reveal two novel interactions between the flexible N- and the C-terminal regions of PrP upon Cu\(^2+\)-binding as follows: (i) between the N-terminal region and helix-1 (residues 144–147) and its nearby loop region (residues 139–143) and (ii) between the N-terminal region (residues 60–91) and helix-2 (residues 174–185) of PrP. We believe these observations have physiological implications.

EXPERIMENTAL PROCEDURES

All chemicals and reagents were of analytical or ultra pure grade. Isopropyl 1-thio-β-D-galactopyranoside was obtained from Bangalore Genei (India), and urea was procured from United States Biochemical (Cleveland, OH), and ampicillin was purchased from Biochem Pharmaceutical Industries Ltd. (India). Protease inhibitor mixture was bought from Roche Diagnostics; NaCl and glycine were from Qualigens, Fine Chemicals (India); CuSO\(_4\) was from Sisco Research Laboratories Pvt. Ltd. (Mumbai, India). Nickel-nitrilotriacetic acid matrix was obtained from Qiagen (GmbH, Hilden, Germany). Other chemicals or reagents were from Sigma. Water used in all reactions was obtained from MilliQ, Millipore (Bedford, MA).

Cloning, Expression, and Purification of Prion Protein Variants—We have cloned, expressed, and purified mouse PrP(23–231) as described earlier (35). PrP(90–231) was cloned into pET21a vector after PCR amplification from PrP(23–231)-pET21a template using gene-specific primers (forward primer TGGCATATGGCCAAAGGGGTAC and reverse primer GCTTTGGAATTCGCTTGATCTCCCG). Inserts in pMOS blue vector (Novagen) were digested with NdeI and HindIII and ligated by T4-DNA polymerase (Promega) in pET21a vector. Escherichia coli DH5α strain was transfected with the plasmid constructs. Positive colonies were screened with the help of ampicillin. The sequence of this construct was verified using 3700ABI automated DNA sequencer. MoPrP(90–231) (pET21a vector) was expressed in E. coli Rosetta DE3. After 12 h of induction, cells were harvested, lysed, and centrifuged. The insoluble pellet was washed and dissolved in 8 M urea and 10 mM reduced glutathione and bound to nickel-nitrilotriacetic acid matrix. On-column oxidative folding of PrP was done by slow removal of the denaturant and the reducing agent. Protein was eluted in 50 mM imidazole. The eluted protein was extensively dialyzed against MilliQ water and then concentrated. Aliquots of protein were stored at −70 °C. The protein concentrations were estimated using the extinction coefficient of 2.70 and 1.70 at 280 nm for 1 mg/ml solutions of PrP(23–231) and PrP(90–231), respectively. The purity of the protein was checked using SDS-PAGE and found to be free of any contaminants.

PrP Preparations for Spectroscopic Measurements—Scattering experiments were performed in 20 mM sodium phosphate buffer (pH 7.4) and 100 mM NaCl, unless otherwise specified. For ITC, SAXS, dynamic light scattering, sedimentation velocity, and NMR experiments, PrP in water with pH 7.4 maintained by dilute NaOH solution was used to avoid aggregation. At low protein (∼0.25 mg/ml) concentrations, the reversible aggregation was not noticeable. However, at higher concentrations up to 2 mg/ml in water (pH 7.4), reversible aggregation was observed below 22 °C.

Cu\(^2+\)-binding Studies—For all Cu\(^2+\)-binding studies, including aggregation, NMR, SAXS, sedimentation velocity, and ITC experiments, Cu\(^2+\) was complexed with glycine in a 1:4 molar ratio to form Cu-(Gly)\(_2\) complex (referred to as Cu\(^2+\) everywhere in the text) to avoid any hydroxide formation. The Cu\(^2+\)(Gly)\(_2\) complex was maintained at pH 6.0 to avoid drastic change in pHe of PrP (water, pH 7.4) upon titration in ITC experiments. After titration, no significant change in pH was observed. Except for ITC, all the binding studies were done using 6 mM eq of Cu\(^2+\).

Scattering Studies—Scattering measurements were monitored using Spectrofluorimeter Fluorolog-3 model FL-3–22 from Horiba Jobin Yvon, Inc. Rayleigh scattering of solutions was recorded at any given temperature with excitation and emission monochromators fixed at 465 nm, and bandpasses for excitation and emission were set at 0.5 and 1 nm, respectively. Data max software provided by the manufacturer was used to monitor scattering. The instrument takes 2.5 and 3 min to increase or decrease the sample temperatures from 15 to 37 °C, respectively. To measure the extent of reversibility, scattering was monitored at various temperatures between 37 and 15 °C and brought back to 37 °C with similar intervals.

Isothermal Calorimetry—Binding of Cu-(Gly)\(_2\) complex with PrP(23–231) was performed using VP-ITC microcalorimeter from MicroCal, LLC. Data acquisition and analysis were done with software provided by the manufacturer. PrP(23–231) at a concentration of 0.25 mg/ml (12 μM) in water (pH 7.4) was used and 2 mM of Cu-(Gly)\(_2\) was titrated against it. Each solution was filtered or centrifuged and degassed before binding experiments. Initial injection of 2 μl followed by 29 injections of 4 μl was used for binding reactions. To remove the heat of dilution generated because of injection of ligands to the protein solution in the binding reactions, base-line correction for raw isotherms was performed by subtracting average saturated values (obtained from six points) from its corresponding isotherms. For analysis, a sequential mode with four binding sites was used. Binding reactions were performed at 16, 20, 25, 30, and 37 °C.
Three reactions were done at each temperature, and averaged values are reported.

**NMR of Prion Protein**—Uniformly $^{15}$N-labeled protein samples of PrP(23–231) and PrP(90–231) were prepared by growing *E. coli* cells in M9 minimal medium containing $^{15}$NH$_4$Cl as the sole source of nitrogen. Protein samples (2 mg/ml) were prepared in ultra pure water ($10^{-18}$ Ω$^{-1}$ cm$^{-1}$ resistivity) whose pH was maintained at 7.4. Two-dimensional $^{15}$N-$^1$H HSQC spectra were recorded for PrP(23–231) and PrP(90–231) at various temperatures ranging from 16 to 37 °C. Each $^{15}$N-$^1$H HSQC was recorded with 1024 and 128 complex points along $^1$H and $^{15}$N dimensions, respectively, using watergate sequence for water saturation, on a Bruker Avance 800 MHz NMR spectrometer equipped with a 5-mm triple-resonance cryogenic probe, whose temperature was calibrated using methanol (280–300 K). All the spectra were referenced using 4,4-dimethyl-4-silapentane-1-sulfonic acid as an external reference at 298 K and then adding the correction of 0.01 ppm/K for the temperature change. The spectra were processed with 4096 and 1024 points along $^1$H and $^{15}$N dimensions, and a window function of 60°-shifted sine square bell was used for apodization along both the dimensions using TopSpin 2.0 software. Analysis was carried out manually using computer-aided resonance assignment (CARA).

**NMR Assignments**—Prior to the assignment of the peaks in the two-dimensional $^{15}$N-$^1$H HSQC of PrP(90–231), recorded in a mixed solvent of $^2$H$_2$O and H$_2$O (pH 7.4), we recorded the same spectrum at pH 7.0 for which the chemical shift information (32) is available in the BioMagResBank (code 16071) under similar buffer conditions, 5 mM phosphate buffer (pH 7.0). All the observed peaks in the pH 7.0 spectrum, totaling 128 out of 138, could be assigned by visual inspection. The missing spectral information about the 10 remaining residues belongs to the flexible part of the protein. In such an assignment, no changes were noticed in the respective chemical shifts, except for the residue Phe-141. We then assigned the $^{15}$N-$^1$H HSQC of PrP(90–231) recorded in water at pH 7.4 by comparing it with the spectral assignments made at pH 7.0. Once again, no spectral changes were noticed except for the absence of 17 more cross-peaks belonging to the unstructured part of the protein (residues 90–120). Thus, we could unambiguously assign the spectral signatures of 111 residues. Such assignments were further used to assign 85 of the observed peaks in the $^{15}$N-$^1$H HSQC of wild-type prion protein, recorded under identical experimental conditions. Among the residues, whose spectral signatures were absent in the spectrum, 28 belong to the unstructured polypeptide stretch (residues 90–120), and the residues Val-166–Asn-171, Arg-230, and Ser-231, all belonging to the globular domain of the protein.

**Sedimentation Velocity Measurements**—Sedimentation velocity measurements were performed using Optima XL-I analytical ultracentrifuge (Beckman Coulter, Fullerton, CA). Prion protein samples (2 mg/ml (pH 7.4), conditions identical to those used in NMR experiments), with and without Cu$^{2+}$, were subjected to centrifugation at 58,000 rpm ($\sim$240,000 × g) at 37 °C, using An60Ti rotor with an aluminum center piece. The protein boundary was scanned at 1-min intervals for 8 h by interference measurements using 30-milliwatt 660 nm laser diode. The sedimentation coefficient $s_{20,w}$ and molecular mass of the protein were calculated using the program SEDFIT (36), which uses nonlinear regression fitting of the sedimenting boundary profile with the Lamm Equation,

$$\frac{dc}{dr} = \frac{1}{r} \times \frac{d}{dr} \left( \frac{Dc}{dr} - s\omega^2 \cdot c \right)$$

which describes the concentration distribution $c(r,t)$ of a species with sedimentation coefficient $s$ and diffusion coefficient $D$ in a sector-shaped volume and in the centrifugal field $\omega^2 r$.

**SAXS Studies**—SAXS measurements were performed with a HECUS S3-Micro SAXS camera attached to a Xenocs microbeam delivery system (copper target, wavelength $\lambda = 1.54$ Å) operating at a power of 50 watts. SAXS measurements of PrP samples with and without Cu$^{2+}$ (2 mg/ml in water (pH 7.4)) at 37 °C for 10,800 s were recorded with a Pilatus 100k pixel detector. The two-dimensional image was reduced to a one-dimensional scattering curve using the Fit2D software and processed using PRIMUS (37). The $R_g$ and $I(0)$ values were obtained using the Guinier approximation (38). To obtain shape information, the scattering data were converted to $P(r)$ function by GNOM software, and the $D_{max}$ was calculated (39). The $P(r)$ function was taken to GASBOR, to generate *ab initio* models (40). This process was repeated 10 times and then averaged using DAMAVER (41). The high resolution NMR data of prion protein and the *ab initio* SAXS envelopes of PrP in the absence and presence of Cu$^{2+}$ obtained from GASBOR program were matched using SUPCOMB (42).

**RESULTS**

**Reversible Aggregation of PrP(23–231) upon Cu$^{2+}$-binding**—We measured aggregation of PrP upon Cu$^{2+}$-binding by monitoring light scattering of the solution under physiological conditions (temperature 37 °C and pH 7.4). Six molar equivalents of Cu$^{2+}$ in the form of Cu-(Gly)$_2$ (see under "Experimental Procedures") were added to full-length PrP (Fig. 1A). No changes in the scattering at 37 °C were observed upon Cu$^{2+}$-binding to PrP(23–231), indicating the absence of aggregation at this temperature (see Fig. 1A, before 200 s). We made a serendipitous observation that as the temperature was lowered to 15 °C, there was a sudden increase in the scattering, suggesting protein aggregation (200–400 s). When the temperature was raised back to 37 °C, the observed light scattering suddenly decreased (400–600 s), indicating reversible protein aggregation. The extent of reversibility was monitored by continuous heating and cooling of the sample for 14 times (supplemental Fig. S2A). The aggregation was reversible except for a minor decrease in the observed light scattering. Temperature dependence of this process (supplemental Figs. S1, A and B, and S2A) suggests that the aggregation occurs below 20 °C. However, at higher concentrations of PrP(23–231), the protein aggregates at higher temperatures (below 25 °C). We also observed that the extent of reversible aggregation increased with increasing concentration of PrP(23–231) (supplemental Fig. S1, C and D). Although this aggregation occurs at a nonphysiological temperature, a detailed understanding of this process is likely to provide insights into the molecular basis of Cu$^{2+}$/PrP interactions.


**Cu²⁺-PrP, Aggregation, and N- and C-terminal Interactions**

**A**

![FIGURE 1. Temperature-dependent reversible aggregation of Cu²⁺-bound PrP. A. reversible aggregation of PrP(23–231) in the presence of Cu²⁺. Gray lines indicate temperature of the sample, and slopes of lines indicate decrease or increase in the temperature. B. reversible aggregation of similar concentrations of PrP(23–231) (circles), Cu²⁺-bound PrP(23–231) (triangles), PrP(90–231) (diamonds), and Cu²⁺-bound PrP(90–231) (squares). The circles, diamonds, and squares are overlapping. Arrows show the directions of experiments. Arrow pointing downward shows cooling, and arrow pointing upward shows heating of sample.](image)

We have therefore investigated the temperature-dependent, reversible aggregation of Cu²⁺-bound PrP.

To examine the role of electrostatic interactions in the Cu²⁺-bound PrP(23–231) aggregation, we monitored the protein aggregation at varying concentrations of salt (sodium chloride, NaCl) (supplemental Fig. S1, E and F). The extent of aggregation increased with an increase in salt concentration, suggesting the involvement of non-electrostatic interactions such as hydrogen bonds (H-bonds) and hydrophobic interactions in the PrP(23–231) aggregation. PrP(23–231) binds Cu²⁺ in the N-terminal region involving four octapeptide repeats (residues 60–91) and at the sites His-96 and His-111 (2–12). To probe the region(s) responsible for temperature-dependent aggregation of Cu²⁺-bound PrP(23–231), we have generated an N-terminal deletion mutant PrP(90–231), which lacks the four octapeptide repeats (region 23–89). Interestingly, Cu²⁺-bound PrP(90–231) (Fig. 1B, squares) did not show any aggregation, whereas the Cu²⁺-bound PrP(23–231) shows reversible aggregation as expected (Fig. 1B, triangles). This result shows that the region encompassing octapeptide repeats is responsible for the temperature-dependent reversible aggregation of Cu²⁺-bound PrP.

We used an amyloid-specific fluorescent probe thioflavine T (43) to further understand the nature of these aggregates (amyloid or amorphous). We did not observe any change in thioflavine T fluorescence with Cu²⁺-bound PrP(23–231) at various temperatures (16, 20, 25, 30, and 37 °C), indicating that the aggregates formed are amorphous in nature (supplemental Fig. S2B). This result rules out any amyloidogenic conversion of PrP(23–231) upon Cu²⁺-binding in the given temperature range.

**Binding Affinity of Cu²⁺ to PrP(23–231) at Various Temperatures**—It is possible that the observed temperature-dependent aggregation could be due to temperature-dependent changes in the affinity of Cu²⁺ for PrP. We determined the affinity of Cu²⁺ for PrP(23–231) at different temperatures using ITC. The apparent dissociation constants ($K_{d(app)}$) thus obtained from the curve fitting of base-line-corrected isotherms (see “Experimental Procedures”) did not show significant differences at various temperatures (statistical analysis shows $p$ values in the range 0.11 to 0.23) (supplemental Fig. S2C). This result clearly suggests that the Cu²⁺-induced reversible aggregation of PrP(23–231) cannot be attributed to changes in the binding affinities.

The ITC data suggest that both $\Delta H$ and $\Delta S$ show linear changes with temperature (Fig. 2, A and B); although $\Delta H$ decreased, $\Delta S$ increased with temperature, indicating conformational changes associated with Cu²⁺-binding to PrP(23–231). Interestingly, we observed a linear correlation between $\Delta H$ and $\Delta S$ with a slope of 297.8 K (Fig. 2C). Such a linear and positive correlation between $\Delta H$ and $\Delta S$, with a slope of ~300 K, has been proposed to indicate enthalpy/entropy compensation for small solutes and proteins in water solutions (44, 45). Temperature-dependent compensation requires heat capacity to remain more or less constant throughout the temperature range used (46). In the present case, the calculated molar heat capacity indeed was found to be constant (~517 cal/mol/K, see Fig. 3D). This observation may be taken to suggest that at high temperatures the process is entropy-driven, whereas at lower temperatures it is driven by enthalpy.

**Novel Interactions between the N- and C-terminal Regions upon Cu²⁺-binding**—Isothermal titration calorimetry studies, *inter alia*, show positive and large heat capacity change upon Cu²⁺-binding to PrP(23–231) (~517 cal/mol/K, see Fig. 2D). Several studies associate positive and large heat capacity change with the breaking of H-bonds or long range electrostatic interactions or loss of polar residues on the surface- or temperature-dependent conformational changes (47). In our experiments, positive and large heat capacity change suggests that PrP(23–231) undergoes conformational changes upon binding to Cu²⁺ at various temperatures. We have investigated the possible structural alteration using NMR and SAXS. Because PrP is a glycosylphosphatidylinositol-anchored protein, it has been suggested by Nishina et al. (48) that it is likely to exist in relatively nonpolar micro-environments with a dielectric constant similar to those of low ionic strength buffer solutions. We have therefore carried out these experiments under low ionic strength conditions (see “Experimental Procedures”). We have recorded a series of $^{15}$N-$^1$H HSQC spectra of PrP(23–231) (full length) at different temperatures in the presence and absence of Cu²⁺ (Fig. 3; supplemental Fig. S3). We could monitor cross-peaks from the C-terminal region of PrP (residues 120–231). The cross-peaks from the N-terminal region (residues 23–120), which included Cu²⁺-binding sites such as the octapeptide...
 repeats region (residues 60–91) and His-96/His-111, were not visible in the given spectra in the presence and absence of \( \text{Cu}^{2+} \) (see “Experimental Procedures”). This facilitated observation of specific conformational changes in the C-terminal region of PrP(23–231) induced by \( \text{Cu}^{2+} \)-binding to the N-terminal region. Being paramagnetic in nature, \( \text{Cu}^{2+} \) is known to increase transverse (spin-spin) relaxation rates, thereby broadening the NMR spectral lines (49). Upon \( \text{Cu}^{2+} \) addition, we observed the line broadening of only some but not all the peaks, indicating strong binding of \( \text{Cu}^{2+} \) to specific sites. Several of the original peaks in the HSQC spectrum of PrP(23–231) broadened, upon addition of \( \text{Cu}^{2+} \), although others remained unaffected in their line widths. The intensities of these broadened peaks were below the NMR detection limit (Fig. 3). We further noted that the number of broadened cross-peaks increased with a decrease in temperature in the HSQC spectra of \( \text{Cu}^{2+} \)-bound PrP(23–231) (compare blue cross-peaks in Fig. 3A with those in B and those in supplemental Fig. S3, A and F). However, in the absence of \( \text{Cu}^{2+} \), no such broadening of cross-peaks was noticed in the HSQC spectra of PrP(23–231) (compare red (without \( \text{Cu}^{2+} \)) and blue (with \( \text{Cu}^{2+} \)) cross-peaks in Fig. 3, A and B). At 37 °C, cross-peaks arising from Phe-141, Asn-143, and Asp-147 were found missing (Fig. 3A; Table 1; and Fig. 4, C, D, and F). As the temperature was lowered, peaks arising from Glu-146 and Ile-139 disappeared at 34 °C (Fig. 4, A and B). We also carried out dynamic light scattering measurements (under conditions identical to NMR experiments) and found no change in the hydrodynamic radius of PrP and \( \text{Cu}^{2+} \)-bound PrP (supplemental Fig. S2D). We also carried out sedimentation velocity measurements with PrP and \( \text{Cu}^{2+} \)-bound PrP at 37 °C. Fig. 6, A and B, shows the movement of sedimenting boundary, represented as fringe displacement with time, of the samples of PrP and \( \text{Cu}^{2+} \)-bound PrP, respectively. Fig. 6C shows the sedimentation coefficient profiles of PrP \( (s_{20, w} = 2.694) \) and \( \text{Cu}^{2+} \)-bound PrP \( (s_{20, w} = 2.862) \) derived from curve-fitting analyses of the sedimentation profiles. Sedimentation profiles indicate that ~98.5% of the population in the samples containing PrP and \( \text{Cu}^{2+} \)-bound PrP are in monomeric form with a molecular mass of ~23.4 kDa. The remaining negligible fraction (~1.5%), which corresponds to higher oligomers (including dimers, trimers, or tetramers), is observed both in PrP and \( \text{Cu}^{2+} \)-bound PrP, clearly showing that addition of \( \text{Cu}^{2+} \) to PrP
samples at 37 °C does not lead to the formation of higher order assemblies. It is important to note that the sedimentation velocity and dynamic light scattering measurements were carried out under the same conditions of concentration of PrP, pH, and temperature as used in the NMR experiments. All these observations taken together clearly show that proximity of paramagnetic Cu²⁺/H₁₁₀₀₁, but not the formation of higher order assemblies, is responsible for the observed line broadening in HSQC spectra.

Interestingly, none of the residues in helix-1 and its nearby loop region (Ile-139–Asp-147) are implicated in Cu²⁺/H₁₁₀₀₁-binding to PrP. It is worth mentioning here that in the polypeptide stretch Ile-139 to Asp-147, His-140, not detected in the absence or presence of Cu²⁺/H₁₁₀₀₁ in the given HSQC spectrum, is the lone residue that could bind Cu²⁺/H₁₁₀₀₁. However, studies by Van Doorslaer and co-workers (33) have clearly ruled out the possibility of Cu²⁺-binding at this residue. Therefore, we believe that the reason for the disappearance of peaks from these regions can only be explained if Cu²⁺ bound at other sites such as the octapeptide repeats region and/or His-96/His-111 of the N-terminal region might become proximal to these residues and thus cause the paramagnetic line broadening leading to the disappearance of their spectral signatures.

### TABLE 1

| Residue | 37 °C | 34 °C | 31 °C | 28 °C |
|---------|------|------|------|------|
| Ala-120 | Ala-120 | Ala-120 | Ala-120 | Ala-120 |
| Phe-141 | Phe-141 | Phe-141 | Phe-141 | Phe-141 |
| Asn-143 | Asn-143 | Asn-143 | Asn-143 | Asn-143 |
| Asp-147 | Asp-147 | Asp-147 | Asp-147 | Asp-147 |
| Glu-146 | Glu-146 | Glu-146 | Glu-146 | Glu-146 |
| Asn-174 | Asn-174 | Asn-174 | Asn-174 | Asn-174 |
| Asp-178 | Asp-178 | Asp-178 | Asp-178 | Asp-178 |
| Ile-139 | Ile-139 | Ile-139 | Ile-139 | Ile-139 |
| Cys-180 | Cys-180 | Cys-180 | Cys-180 | Cys-180 |
| Asn-181 | Asn-181 | Asn-181 | Asn-181 | Asn-181 |
| Cys-170 | Cys-170 | Cys-170 | Cys-170 | Cys-170 |
| Ile-182 | Ile-182 | Ile-182 | Ile-182 | Ile-182 |
| Lys-185 | Lys-185 | Lys-185 | Lys-185 | Lys-185 |
| Arg-208 | Arg-208 | Arg-208 | Arg-208 | Arg-208 |

a Ala-120 is present in a flexible N-terminal region, and Arg-208 is present in helix-3.

b In the case of PrP(90–231), the cross-peak arising from Ile-139 was found to disappear at 25 °C.

### FIGURE 3

Two-dimensional ¹⁵N-¹H HSQC spectra of PrP(23–231) and PrP(90–231). A and B, two-dimensional ¹⁵N-¹H HSQC spectra of PrP(23–231) (red cross-peaks) were overlaid with those of Cu²⁺-bound PrP(23–231) (blue cross-peaks) at 37 °C (A) and 25 °C (B). C and D, two-dimensional ¹⁵N-¹H HSQC spectra of PrP(90–231) (red cross-peaks) and Cu²⁺-bound PrP(90–231) (blue cross-peaks) were at 37 °C (C) and 25 °C (D).
To probe residues from the N-terminal regions that might interact with helix-1 and helix-2 of the C-terminal region, we have recorded $^{15}$N-$^1$H HSQC spectra of the N-terminal deletion mutant PrP(90–231), which lacks the four octapeptide repeats (region 23–89) leaving His-96/His-111, as the only Cu$^{2+}$-binding site. We noticed that the cross-peaks arising from Phe-141, Asn-143, Glu-146, Asp-147, and Asp-178 completely disappeared upon addition of Cu$^{2+}$ at 37 °C (Figs. 3C–4G). Upon lowering the temperature to 25 °C, the cross-peak arising from Ile-139 also disappeared (Figs. 3D and 4B; supplemental Fig. S4B; in Table 1 see the boldface residues). Interestingly, the cross-peaks of PrP(90–231) from helix-1 and its nearby loop region (Ile-139–Asp-147), which broadened in the presence of Cu$^{2+}$ at 37 °C, were similar to those of PrP(23–231), as mentioned above. PrP(90–231) did not show protein aggregation in the presence of Cu$^{2+}$ at low temperatures (Fig. 1B). We therefore propose here that the C-terminal region (Ile-139 to Asp-147) of PrP is involved in a novel interaction with the Cu$^{2+}$-bound N-terminal region (residues 90–120 that contains the site His-96/His-111). The role of the octapeptide repeats in this interaction is less pronounced as Cu$^{2+}$-bound PrP(90–231) does not have residues from 23 to 89 that harbor octapeptide repeats. Thus, on the basis of our observations in proteins PrP(90–231) and PrP(23–231), we suggest that the region 90–120 (which contains the site His-96/His-111) becomes proximal to the region Ile-139–Asp-147 for interaction, upon binding to Cu$^{2+}$.

PrP(23–231) also shows broadened cross-peaks from the helix-2 region, which is not observed in the case of PrP(90–231). A few reports suggest that Cu$^{2+}$ binds to His-187 (9–12). However, we found that the cross-peak arising from His-187 broadened only below 22 °C in the HSQC spectra of Cu$^{2+}$-bound PrP(23–231) but not in that of Cu$^{2+}$-bound PrP(90–231) (Table S1). Because both the variants of PrP (PrP(23–231) and PrP(90–231)) have His-187, they should have shown a similar broadening phenomenon if Cu$^{2+}$ binds to this site. Moreover, the broadened cross-peaks arising from the helix-2 region (residues 174–182 until 28 °C) are distal to His-187. Thus, the NMR data clearly shows that His-187 is not involved in Cu$^{2+}$-binding under the chosen experimental conditions. This observation indicates the presence of Cu$^{2+}$ in close proximity of the helix-2 region, which might be responsible for the observed line broadening at high temperatures.
broadening. We believe that the reason for the disappearance of peaks from this region can only be explained by a second novel interaction of the N-terminal domain with the helix-2 region. In the case of PrP(90–231) (which lacks octapeptide repeats), we did not observe broadening of peaks in the helix-2 region suggesting the role of Cu\textsuperscript{2+}/H11001-bound octapeptide repeats in the disappearance of cross-peaks from the helix-2 region. Therefore, we further propose that Cu\textsuperscript{2+}-bound octapeptide repeats region (residues 60–91) interacts with the helix-2 region (residues 174–185) and is responsible for the broadening of cross-peaks arising from this region. Viles et al. (28) investigated a small region of PrP covering 51–91 residues that harbor the Cu\textsuperscript{2+}-binding octapeptide repeats region and observed line broadening of some lines from this region. Because we have used full-length PrP(23–231), we were able to observe long range, intra-protein interactions upon Cu\textsuperscript{2+}-binding.

Our NMR results also show that residues present in hydrophobic core were less affected until 24 °C. Upon further lowering the temperature to 22 °C and below, Cu\textsuperscript{2+}-bound PrP(23–231) exhibited protein aggregation. HSQC spectra recorded at these temperatures showed general broadening of cross-peaks originating from residues of helices 1 and 2 (supplemental Table S1) and a few of the residues belonging to helix-3, which are in contact with these helices. As described earlier, the octapeptide repeats are responsible for the temperature-dependent reversible aggregation of Cu\textsuperscript{2+}-bound PrP(23–231). On the basis of NMR results, we suggest the role of interaction of helix-2 and octapeptide repeats region in the temperature-dependent reversible aggregation of Cu\textsuperscript{2+}-bound PrP(23–231).

Compaction of the Extended Conformation of PrP upon Cu\textsuperscript{2+}-binding—Our results indicate that the line broadening of the peaks from helix-1 and helix-2 regions above 24 °C is not due to the formation of higher order assemblies but a result of the interaction of helix-2 and octapeptide repeats region in the temperature-dependent reversible aggregation of Cu\textsuperscript{2+}-bound PrP(23–231). To get an insight into these intra-molecular interactions of the N-terminal region with the C-terminal counterpart, we estimated \( R_g \) and \( D_{max} \) of PrP(23–231) and Cu\textsuperscript{2+}-bound...
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PrP(23–231) using SAXS. The $R_g$ of PrP(23–231) (2.55 nm) and Cu Superscript 2+-bound PrP(23–231) (2.46 nm) determined by Guinier analysis show small differences (left inset, Fig. 7A). Distance distribution function, $P(r)$ analysis, reveals reduction in the $D_{max}$ value from 10 nm for PrP(23–231) to 9 nm for Cu Superscript 2+-bound PrP(23–231) (Fig. 7B), which suggests that Cu Superscript 2+-bound PrP(23–231) might have become more compact compared with PrP. The N-terminal region of PrP is unstructured and flexible in nature. Hence, we analyzed the data using ensemble optimization method (EOM), which gives useful information such as $R_g$ and $D_{max}$ distributions in the case of proteins with flexible domains. Fig. 7, C and D, compare the $D_{max}$ and $R_g$ distributions, respectively, of the ensembles of the selected conformations with the $D_{max}$ and $R_g$ distributions of the initial pool (randomly generated) of 10,000 structures. The $D_{max}$ and $R_g$ distributions of PrP(23–231) and Cu Superscript 2+-bound PrP(23–231) showed multimodal distributions. To discern these distribution curves, we have deconvoluted the $D_{max}$ and the $R_g$ distribution plots (supplemental Fig. S5, A–D). In the $D_{max}$ distribution of PrP(23–231), the two conformer populations (mean around 7.31 and 9.52 nm with fractional contributions of 48.7 and 49.9%, respectively) (Fig. 7C, red trace; supplemental Fig. S5B) converged into a major population with $D_{max}$ 8.39 nm (72.7%) upon Cu Superscript 2+-binding (Fig. 7C, blue trace; supplemental Fig. S5D). This suggests that a significant amount of compaction of the extended conformation of PrP(23–231) occurs upon Cu Superscript 2+-binding. We also found similar features in the $R_g$ distributions of PrP(23–231) and Cu Superscript 2+-bound PrP(23–231). $R_g$ distributions of PrP(23–231) corresponding to 2.25 nm (46.4%) and 3.05 nm (34.5%) (Fig. 7D, red trace; supplemental Fig. S5A) are changed to 2.24 nm (22.8%) and 2.65 nm (73.4%) (Fig. 7D, blue trace; supplemental Fig. S5C), respectively, upon Cu Superscript 2+-binding, which supports the $D_{max}$ distribution data obtained above. The results from the EOM analysis of the size distribution are in agreement with the values obtained from $P(r)$ distribution function, where reduction of 1 nm is seen for $D_{max}$ of Cu Superscript 2+-bound PrP(23–231) compared with that of PrP(23–231). The reduction in the $D_{max}$ leading to the global compactness of Cu Superscript 2+-bound PrP(23–231) might be the result of a decrease in the flexibility of the N-terminal region, which exhibits interaction with the C-terminal region of PrP(23–231) upon Cu Superscript 2+-binding. Study from Viles et al. (28) suggests that upon Cu Superscript 2+-binding the octapeptide repeats region achieves compact structure due to four coordinating histidine residues. Studies from Takeuchi et al. (52) have reported that Cu Superscript 2+ induces helix formation in the octapeptide repeat region (53). However, in both these studies short peptides comprising octapeptide or the repeat region (residues 51–91) have been used. We report here a global compaction of full-length PrP(23–231) due to inter-domain interactions upon Cu Superscript 2+-binding. Thus, our SAXS results lend credence to the existence of long range interactions between the N- and C-terminal regions in the Cu Superscript 2+-bound PrP(23–231). The observed marginal increase (Fig. 6C) in the sedimentation coefficient of Cu Superscript 2+-bound PrP (2.86 from 2.69 for PrP) also indicates global compaction.

**DISCUSSION**

We have observed temperature-dependent reversible aggregation of Cu Superscript 2+-bound PrP(23–231); it aggregates at lower temperatures and resolubilizes at physiological temperature. We believe that a novel interaction between helix-2 and the octapeptide repeats region of PrP(23–231) is involved in this reversible aggregation. Our study further suggests that the aggregation might proceed through the formation of hydrogen bonds (H-bonds), which might dissociate rendering the PrP(23–231) soluble at physiological temperature. We have investigated the conformational consequences of Cu Superscript 2+-binding to PrP(23–231) under physiological conditions. On the basis of our results, we propose novel interactions between the N- and C-terminal regions involving the flexible region and the helices-1 and -2. Aggregation of Cu Superscript 2+-bound PrP—Cu Superscript 2+ is known to play a role in prion diseases. It is shown to promote and prevent the process. We noticed that most of the Cu Superscript 2+-binding experiments are either performed at nonphysiological temperatures (2, 10, 13–17) or at very high concentrations of Cu Superscript 2+ (54, 55). Under physiological conditions, we find that Cu Superscript 2+-bound PrP(23–231) does not aggregate. Interestingly, it starts to aggregate when the temperature is lowered. Aggregation at lower temperature has been reported earlier (2, 10, 13–17). Our results suggest that Cu Superscript 2+ may not play a role in the aggregation of PrP at physiological temperature (37 °C). Our experiments with mutant protein (PrP(90–231)) show that the region...
encompassing octapeptide repeats (residues 60–91) is responsible for the temperature-dependent reversible aggregation of Cu²⁺-bound PrP(23–231) (Fig. 1). Our ITC experiments show that the observed temperature-dependent aggregation might not be due to changes in the affinity of Cu²⁺-binding. Binding affinity remains invariant; however, enthalpy and entropy changes show linear and reciprocal relation indicating an enthalpy-entropy-compensation phenomenon (44, 45). The observed enthalpy/entropy compensation phenomenon might be useful in understanding the temperature-dependent aggregation; due to higher entropy at 37 °C, the Cu²⁺-binding sites will be flexible and solvated and might disrupt any cohesive inter-molecular interactions. This facilitates Cu²⁺-bound PrP(23–231) to remain in solution at 37 °C. However, at lower temperatures, these sites might come together initiating cohesive interactions via H-bonds or hydrophobic interactions. Hydrophobic interactions are known to increase in strength with increasing temperatures (56). As mentioned earlier, we observed that Cu²⁺-bound PrP(23–231) aggregates at low temperature, but not at higher temperature (supplemental Figs. S1, A and B, and S2A), indicating a rather lesser role for hydrophobic interactions in the process. Thus, we believe that H-bonds might play a prominent role in the temperature-dependent aggregation of Cu²⁺-bound PrP(23–231).

**Interaction between the N- and C-terminal Regions**—The possibility of interaction between the N- and C-terminal regions of PrP(23–231) has been indicated in earlier studies. Binding of antibodies specific to the N- and C-terminal regions (57), high pressure (58), and NMR studies indicates the presence of a transient interaction of the C-terminal region of helix-2 (residues 187–193) and helix-3 (residues 217–226) with the unfolded N-terminal domain (34). Our Cu²⁺-binding studies using ITC revealed changes in enthalpy with temperatures that resulted in large positive heat capacity change. Large heat capacity change reflects structural rearrangements upon Cu²⁺-binding to PrP(23–231). Our NMR studies indicate previously unknown interactions of the N-terminal region with the C-terminal region in the Cu²⁺-bound PrP(23–231). The NMR of full-length PrP has been reported earlier; there are also reports of NMR studies on Cu²⁺-binding to fragments of PrP (28, 32, 59). To the best of our knowledge, this is the first investigation of NMR of Cu²⁺-bound full-length PrP, which shows novel long range inter-domain interactions.

**Fig. 8** schematically describes temperature-dependent reversible aggregation and long range inter-domain interactions observed with Cu²⁺-bound PrP(23–231). PrP (Fig. 8, panel 1) shows interaction of the N-terminal region with the C-terminal region at helix-2 and -3 (residues 187–193 and 217–226, respectively) (green arrow) as reported earlier (55–57). At 37 °C, upon Cu²⁺ addition, our studies suggest interaction of the N-terminal flexible domain with helix-1 and its nearby loop region (Fig. 8, panel 2, blue arrow). To the best of our knowledge, this is the first report of the interaction of helix-1 and its nearby region (residues 139–147) with the flexible N-terminal...
region (residues 90–120 that contains the site His-96/His-111). Cu²⁺-bound PrP(23–231), in the temperature range 37–24 °C, exhibits another novel interaction between the N-terminal (residues 60–91) and the C-terminal (residues 174–185) regions in the presence of Cu²⁺ (Fig. 8, panel 3, red arrow). Aggregation at lower temperatures might be initiated via interaction of helix-2 and the Cu²⁺-bound octapeptide repeats region. This interaction will further result in aggregation of Cu²⁺-bound PrP (Fig. 8, panel 4).

Our SAXS results lend credence to the scheme described in Fig. 8. We have generated low resolution ab initio SAXS models for PrP(23–231) and Cu²⁺-bound PrP(23–231) (supplemental Fig. S6). Cu²⁺-bound PrP(23–231) (orange envelope) shows more curved and smaller envelope compared with that of PrP(23–231) (blue envelope) (supplemental Fig. S6, A and B). It is possible that the interaction of the flexible N-terminal region with the C-terminal region, upon Cu²⁺-binding, leads to the observed changes in the shape of envelope of Cu²⁺-bound PrP(23–231) compared with that of PrP(23–231).

**Functional Consequence of Interaction of Helix-1 and N-terminal Region (Residues 90–120)—** We believe that the interaction of the N-terminal region (residues 90–120 that contain the site His-96/His-111) with the C-terminal region (residues 139–147) is important because of the role played by helix-1 and its nearby loop in the conversion of PrP⊙ to PrP⊙⊙. Involvement of helix-1 in the conversion of PrP⊙ to PrP⊙⊙ was first predicted by Morrissey and Shakhnovich(60) using CHARMM energy calculation. The β-nucleation model proposed by them states that PrP⊙⊙ is an aggregate with a hydrophilic core, consisting of a β-sheet-like arrangement of constituent helix-1 components (60). Helix-1 has been shown to promote aggregation (61, 62). It was also found in the dimer interface of the crystal structure of human PrP (63). Site 139–141 has been shown to be the most pressure-sensitive region (58) and found to have a much higher aggregation tendency compared with helix-1 (64). The helix-1 region Asn-143–Glu-146 also plays a prominent role in the prion conversion in vivo (65). Thus, it is evident that helix-1 and the nearby loop region play a major role in the aggregation and conversion of PrP to PrP⊙⊙. Our study shows that helix-1 and the nearby loop region interact with the N-terminal region (residues 90–120 that contain the site His-96/His-111) upon Cu²⁺-binding. As mentioned earlier, Cu²⁺ inhibits amyloid formation of PrP in vitro (22). Baskakov and co-workers (22) attributed this inhibition to binding of Cu²⁺ in the C-terminal region of PrP. However, even in the absence of the C-terminal region, PrP(82–146), in the presence of Cu²⁺, did not show amyloid fibril formation (23). It is possible that the interaction of the N-terminal region (residues 90–120) with the loop (Ile-139–Phe-141) and helix-1 (Asn-143–Arg-148) region of PrP(23–231) upon Cu²⁺-binding is responsible for the observed inhibition of the amyloid fibril formation in all the three cases, namely PrP(82–146) (23), PrP(89–231) (22), and PrP(23–231) (22). This interaction is not present in PrP(23–231) in the absence of Cu²⁺. We therefore propose an inhibitory role of Cu²⁺ on the conversion of PrP⊙ to PrP⊙⊙ in vitro and in vivo via the interaction between the N-terminal region and helix-1 and its nearby loop region.

**Functional Consequence of Interaction of Helix-2 and Octapeptide Repeats—** PrP(23–231), in contrast to PrP(90–231), undergoes temperature-dependent reversible aggregation upon Cu²⁺-binding possibly because of the interaction between the octapeptide repeats region (residues 60–91) and helix-2. Reversible aggregation of Cu²⁺-bound PrP(23–231) is only observed at lower temperature but not at 37 °C. Therefore, we believe that interaction between the octapeptide repeats region and the helix-2 region primes Cu²⁺-bound PrP(23–231) for inter-protein interactions at 37 °C. Such interactions might have physiological significance.

The globular C-terminal domain of PrP includes two subdomains, H1-B1 and B2-H2-H3. NMR relaxation dynamics show that H1-B1 and the connecting loop are more flexible compared with the B2-H2-H3 subdomain (66). Rare, large scale motions of the subdomains have been proposed to initiate prion protein aggregation (67). Recent studies have shown that separation of these subdomains and formation of domain-swapped dimers are a prerequisite for the aggregation of prion...
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protein (68, 69). This implies that increased proximity between these subdomains might have an opposite effect and might delay or completely abrogate the aggregation process. In this study, we show that N-terminal region wraps around C-terminal region by interacting with helix-1 and helix-2 upon copper binding. This might lead to increased proximity of the subdomains, thus inhibiting the aggregation process. It appears that copper inhibits amyloid formation either by interfering with the region comprising H1 and the nearby loop (see above) or by decreasing separation between the two subdomains of the C-terminal region. These implications are important in the context of amyloid formation and role of copper in prion disease.

Harris and Pauly (70) proposed that conformational changes in PrP⁷C upon Cu²⁺-binding might increase the affinity of the protein for a putative endocytic receptor that localizes PrP⁷C in clathrin-coated pits. Based on our observations, we propose that the interaction between the octapeptide repeats region and the nearby loop in the C-terminal region might be responsible for the conformational changes required for increased endocytosis.

The role of Cu²⁺ as an attenuator or facilitator in prion diseases is highly controversial. Our observation that PrP(23–231) in the presence of Cu²⁺ does not aggregate at 37 °C supports the suggestion that Cu²⁺ attenuates prion disease. However, we have also observed an unusual temperature-dependent aggregation of Cu²⁺-bound PrP(23–231). It aggregated at lower temperature and resolubilized at physiological temperature. We have investigated this phenomenon using various biophysical techniques such as ITC, NMR, and SAXS thus providing detailed thermodynamic and conformational aspects of Cu²⁺-binding to PrP(23–231). We observed two novel long range inter-domain interactions between the N- and C-terminal regions. Our study shows that Cu²⁺-binding His-96/His-111 becomes proximal to helix-1 and its nearby loop region (resi- dues 139–147). This region is shown to have a prominent role in amyloidogenesis in vitro (22) and prion conversion in vivo (23). Our observation suggests that upon Cu²⁺-binding, the proximity of His-96/His-111 to Ile-139–Asp-147 and proximity between the two subdomains of the C-terminal region are involved in the inhibition of amyloid formation. In addition, we also observed interaction of the octapeptide repeats region (residues 60–91) with helix-2 (residues 174–185) upon Cu²⁺-binding. We speculate the involvement of this interaction in the aggregation process at lower temperatures. We believe that these interactions should prove useful in understanding the biological functions of prion protein upon Cu²⁺-binding in vivo such as conformational changes (71), endocytosis (70), lipid rafts localization, and Cu²⁺-induced signal transduction (72, 73) and in developing therapeutic strategies against prion diseases.

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