Calprotectin influences the aggregation of metal-free and metal-bound amyloid-β by direct interaction.
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Proteins from the S100 family perform numerous functions and may contribute to Alzheimer’s disease (AD). Herein, we report the effects of S100A8/S100A9 heterooligomer calprotectin (CP) and the S100B homodimer on metal-free and metal-bound Aβ40 and Aβ42 aggregation in vitro. Studies performed with CP-Ser [S100A8(C42S)/S100A9(C3S) oligomer] indicate that the protein influences the aggregation profile for Aβ40 in both the absence and presence of metal ions [i.e., Cu(II) and Zn(II)]. Moreover, the detection of Aβ40-CP-Ser complexes by mass spectrometry suggests a direct interaction as a possible mechanism for the involvement of CP in Aβ aggregation. Although the interaction of CP-Ser with Aβ40 impacts Aβ40 aggregation in vitro, the protein is not shown to attenuate Aβ-induced toxicity in SH-SYSY cells. In contrast, S100B has a slight effect on the aggregation of Aβ. Overall, this work supports a potential association of CP with Aβ in the absence and presence of metal ions in AD.

Significance to Metallomics
The metallome plays a central role in many human diseases, including Alzheimer’s disease (AD). Metal ion dyshomeostasis and amyloid-β (Aβ) aggregation are implicated in this debilitating neurodegenerative disorder; however, few studies have addressed interactions between transition metal ions, Aβ, and metal-binding proteins that are present in the brain. Thus, possible interactions of Cu(II) and Zn(II), two metal ions central to AD pathology, with Aβ and calprotectin (CP) and S100B were evaluated, and CP was found to noticeably affect Aβ aggregation. The results highlight that the interplay of metal ions, Aβ, and CP may have a consequence for AD pathogenesis.

Introduction
Alzheimer’s disease (AD) is the most prevalent form of dementia and accounts for approximately 60% of neurodegenerative disorders.1-10 This illness begins with mild cognitive impairment, which progresses to severe dementia and ultimately death.7-10 Associated with these cognitive symptoms are histopathological features of AD-affected brains, notably the presence of senile plaques which primarily consist of amyloid-β (Aβ) aggregates.3-10 Aβ is produced in the brain as a result of the proteolytic cleavage of the amyloid precursor protein (APP) by β- and γ-secretases.6-10 A spectrum of differently sized Aβ peptides is generated, ranging from 38 to 43 amino acid residues, but the predominant forms are Aβ39 (ca. 90%) and Aβ42 (ca. 10%).6-14 Both peptides tend to generate oligomeric species that have been implicated as toxic agents, and can aggregate further into mature fibrils that accumulate in senile plaques.7-14 Although both peptides form fibrils, plaques are enriched in Aβ42 compared to Aβ40.15-18

Numerous chemical, biological, and biomedical research initiatives regarding AD pathogenesis have focused on Aβ; however, AD etiology remains ambiguous and other factors are almost certainly involved. This assessment is especially likely considering that therapeutics designed to principally target Aβ have failed to proceed from clinical trials into medical use,16,17 although promising results were indicated in AD mouse models.18,19 Therefore, current research seeks to further elucidate the etiology of AD, which may arise from multiple agents, possibly occurring in tandem and thus underlying the multifactorial nature of the disease. Along with Aβ aggregation, chronic inflammation, elevated oxidative stress, and metal ion dyshomeostasis are pervasive in the brain of AD patients and could contribute to disease onset and progression.8-14 Since multiple parameters in addition to Aβ have been suggested to be involved in AD pathogenesis, much effort has been expended to investigate the interactions of Aβ with other agents present in the brain that may also be linked to inflammation and metal ion dyshomeostasis.11,12,20-23 Among these candidates is the S100 protein family, members of which exhibit diverse functions, including participation in
the inflammatory response, calcium ion homeostasis, and interactions with transition metal ions.\textsuperscript{24–29} Moreover, S100 proteins are implicated as damage-associated molecular pattern molecules (DAMPs), and are thus indicators of a sustained, chronic noninfectious inflammatory response similar to that observed in AD.\textsuperscript{26,30,31}

S100A9 and S100B are among the most widely studied S100 proteins in AD, and both proteins have been found to be elevated in human AD brain tissue as well as AD mouse model brain tissue.\textsuperscript{22,32–34} Knockdown of S100A9 was observed to improve memory deficits in an AD mouse model and lower plaque burden.\textsuperscript{35} S100B ablation in AD-affected mice could similarly attenuate plaque deposition in specific brain regions,\textsuperscript{36} whereas S100B overexpression in the AD mouse model exacerbated plaque deposition and AD pathology.\textsuperscript{37} These studies support possible roles for S100A9 and S100B in AD pathogenesis.

Calprotectin (CP), a heteroooligomer of S100A8 and S100A9 that chelates transition metal ions and participates in the metal-withholding innate immune response,\textsuperscript{38} is relatively understudied in AD research. An \textit{in vitro} study demonstrated that S100A9 could promote Aβ fibrilization,\textsuperscript{21} and \textit{ex vivo} analysis of postmortem AD tissue exhibited elevated levels of aggregated S100A9, possibly co-localized with Aβ, but not S100A8 and thus not CP.\textsuperscript{22,32,34} Nevertheless, this observation does not preclude the possibility that CP and Aβ interact. Recently, there has been growing consensus that structured S100A9 oligomers are relatively more toxic than Aβ fibrils, and a biomolecule that does not aggravate Aβ fibrillization could still exhibit toxicity in AD via structured soluble oligomers.\textsuperscript{39–42} Therefore, CP was selected as a case study and possible S100 protein candidate that may not exacerbate Aβ fibrilization since it was not found to be co-aggregated and increased in AD plaques but may influence Aβ aggregation. The effect of S100B, another metal-binding S100 protein that exists in the brain,\textsuperscript{43–45} on Aβ aggregation was also investigated.

Herein, we report the impact of CP and S100B on the formation of (i) Aβ fibrils and (ii) Aβ oligomers, as well as (iii) the relative distribution of various-sized peptide species of both metal-free and metal-associated Aβ (metal–Aβ). In addition, we demonstrate that CP directly interacts with Aβ by employing electrospray ionization mass spectrometry (ESI-MS). We further establish that the combined presence of both Aβ and CP is not able to improve Aβ-induced cytotoxicity. Although the distribution of S100A8 and S100A9 has been determined in AD mouse model brain tissue,\textsuperscript{35} to the best of our knowledge, this study is the first report on the influence of CP on the behavior (e.g., aggregation and toxicity) of both Aβ\textsubscript{40} and Aβ\textsubscript{42} in the absence and presence of metal ions.

Results and discussion

Selection and preparation of S100 proteins

All apo S100 proteins employed in this work were obtained by recombinant expression in \textit{Escherichia coli}.\textsuperscript{29,46} CP variants were purified as apo heterodimers. The 24-kDa CP heterodimer exhibits two transition-metal-binding sites at the S100A8/S100A9 interface. Site 1 is a His\textsubscript{3}Asp motif comprised of (A8)His83, (A8)His87, (A9)His20, and (A9)Asp30. Site 2 is a His\textsubscript{2} site comprised of (A8)His17, (A8)His27, (A9)His91, (A9)His95, (A9)His103, and (A9)His105. Both of these sites coordinate Cu(II) and Zn(II) with high affinity.\textsuperscript{29,46} In this work, we employed two CP variants, CP-Ser and CP-Ser\textsubscript{A9}, that were biochemically evaluated.\textsuperscript{29} Each S100 subunit of CP contains one Cys residue, and CP-Ser is the S100A8(C42S)/S100A9(C3S) variant. Neither Cys residue is in close proximity to the metal-binding sites, and CP-Ser has been used extensively in order to avoid the use of reducing agents or the formation of disulfide bonds during metal-binding studies.\textsuperscript{29} CP-Ser\textsubscript{A9} harbors eight additional point mutations (His\textsubscript{Asp} → Ala) and cannot bind transition metal ions at the His\textsubscript{Asp} and His\textsubscript{Cys} sites. This protein allowed us to investigate whether the metal-binding capacity of CP influences its interactions with Aβ in the absence and presence of metal ions.

The 21-kDa S100B homodimer also contains two transition-metal-binding sites, which are located at the homodimer interface. At pH 6.5, these sites are defined by His15 and His25 from one monomer, and His85 and Glu89 from the other monomer.\textsuperscript{47,48} Both of these sites coordinate Zn(II) and Cu(II).\textsuperscript{47} In this work, we employed S100B and the S100B\textsubscript{C} variant, which lacks the metal-binding sites (Supporting Information and Fig. S1, ESI†).

Influence of CP-Ser on the formation of β-sheet-rich Aβ aggregates with and without metal ions

The generation of β-sheet-rich aggregates of metal-free Aβ (prepared in Chelex-treated buffer) and metal–Aβ 0.1 or 1.0 equiv of Zn(II) or Cu(II) relative to Aβ) in the absence and presence of CP-Ser and CP-Ser\textsubscript{A9} was monitored by the thioflavin-T (ThT) assay and transmission electron microscopy (TEM) (Fig. 1 and 2 and Fig. S2–S4, ESI†). We first investigated the effect of these proteins on the aggregation of Aβ\textsubscript{40}. Under metal-free conditions, incubation of Aβ\textsubscript{40} with CP-Ser or CP-Ser\textsubscript{A9} led to lower fluorescent responses from ThT than the sample containing Aβ\textsubscript{40} only. ThT becomes fluorescent upon binding to β-sheet-rich aggregates, and its fluorescence intensity is a quantifiable measure of the amount of β-sheet-rich structures present.\textsuperscript{46} Upon treatment of Aβ\textsubscript{40} with CP-Ser or CP-Ser\textsubscript{A9}, a decrease in fluorescence intensity greater than 80% was observed (Fig. 1b). In addition, the aggregation kinetics of Aβ\textsubscript{40} with and without CP-Ser were monitored by the ThT assay (Fig. S2, ESI†). The elongation phase of Aβ\textsubscript{40} aggregation with and without CP-Ser occurred from the beginning of the incubation and reached a plateau after ca. 8 h. The fluorescence intensity of ThT in the samples containing CP-Ser and Aβ\textsubscript{40} was ca. 20% of that observed for the samples of Aβ\textsubscript{40} without CP-Ser after 24 h incubation. This indicates that the formation of a smaller amount of β-sheet-rich Aβ\textsubscript{40} aggregates upon treatment of CP-Ser. Moreover, CP-Ser was
indicated to delay the aggregation kinetics of Aβ40 following the incubation time up to 24 h (Fig. S2, ESI†). Together, CP-Ser was observed to modulate the aggregation of Aβ40.

In the presence of metal ions [i.e., Zn(II) and Cu(II)], the ThT assay indicated that both CP-Ser and CP-Ser∆∆ could influence metal–Aβ fibrillation depending on the equivalents of metal present. CP-Ser reduced the formation of β-sheet-rich Aβ40 aggregates with 1 and 2 equiv of Zn(II) relative to the protein (concentrations of Aβ and CP are equivalent), whereas this protein did not affect aggregation when > 2 equiv of Zn(II) were added (Fig. 1b and Fig. S3b, ESI†). On the other hand, CP-Ser∆∆ may have some influence at sub-stoichiometric levels of Zn(II), but may not have a significant effect on Aβ40 aggregation at supra-stoichiometric Zn(II) (Fig. 1b and Fig. S3b, ESI†). Similar trends were observed upon the addition of Cu(II) to CP-Ser, showing alteration of Aβ40 aggregate formation with up to 1 equiv of Cu(II). In contrast, CP-Ser∆∆ exhibited a marginal effect on Cu(II)–Aβ40 aggregation (Fig. 1b). These results suggest that the presence of CP-Ser or CP-Ser∆∆ reduces the formation of β-sheet-rich (ThT-reactive) Aβ aggregates.50

The ability of CP-Ser to alter Aβ40 aggregation at stoichiometric Zn(II) or Cu(II) (Fig. 1b) compared to CP-Ser∆∆, which had no influence against Aβ40 fibrillation under these conditions, indicates that the metal-binding sites of CP, and hence metal chelation, contribute to the modification of metal–Aβ40 aggregation. The effect of both CP-Ser and CP-Ser∆∆ to impact metal-free Aβ40, which likely adopts distinct structural conformations to metal–Aβ40,52–57 also highlights the importance of protein-protein interactions and structures on the ability of proteins to alter Aβ aggregation, in addition to metal chelation.

Apparent dissociation constant (Kd) values obtained by fluorescence methods have been reported for both CP and Aβ for metal ions. Although these experiments have been performed under different conditions, a comparison of these available Kd values suggests that CP-Ser can disrupt metal binding to Aβ. For instance, the two Zn(II) sites of the CP heterodimer are reported to bind Zn(II) at pH 7.5 with Kd values of 133 pM and 185 nM, by the site 1 and site 2, respectively.58 Recently, CP was reported to bind Cu ion with Kd values of 0.2 pM and 4 pM at pH 7.0, respectively, by the site 1 and site 2.59 On the other hand, reported Kd values obtained by fluorescence measurements were varied for Aβ (for Zn(II), from 1 to 300 µM; for Cu(II), ca. 0.1 nM).60–65 Note that metal binding affinities of Aβ have been shown to be dependent on the technique employed (e.g., isothermal titration calorimetric analysis, UV-vis spectroscopy, and nuclear magnetic resonance spectroscopy) and experimental conditions (e.g., concentrations, temperature, and buffer) [for example, ca. 1–20 µM for Zn(II) and 0.4–70 nM for Cu(II)].64–72 Thus, regardless of the method of analysis and experimental conditions, the data from previous studies indicate that CP-Ser binds Zn(II) and Cu(II) with higher affinities than Aβ.58–72

To probe whether CP-Ser interferes with metal binding to Aβ, competition experiments were performed between Aβ40.
Examination of how S100B and S100B\(_\alpha\) affect A\(\beta\) aggregation revealed the behavior that differs from CP-Ser and CP-Ser\(_\alpha\). Under metal-free conditions, initial inspection by the ThT assay indicated that S100B and S100B\(_\alpha\) could slightly influence metal-free A\(\beta\)\(_{40}\) aggregation because a small attenuation of ThT fluorescence occurred (Fig. 1b). The TEM studies also demonstrated that S100B and S100B\(_\alpha\) could induce small and subtle morphological changes in metal-free A\(\beta\)\(_{40}\) fibrilization, resulting in the formation of shorter metal-free A\(\beta\)\(_{40}\) fibrils with fewer amorphous aggregates, compared to the sample of A\(\beta\)\(_{40}\) only (Fig. 2b). As the morphologies of both S100B and S100B\(_\alpha\) aggregates were noticeably different from A\(\beta\) aggregates, we could distinguish accumulated A\(\beta\) and S100B/S100B\(_\alpha\) species (Fig. 2b). The truncated fibrils that form in the presence of S100B, relative to the broader and longer fibrils in the samples of A\(\beta\)\(_{40}\) only, bind fewer ThT molecules and hence generate a slightly lower fluorescence signal in the ThT assay (Fig. 1b). Since fibrils are still formed, the fluorescence signal is not as effectively abrogated as observed for CP-Ser with A\(\beta\)\(_{40}\).

In the presence of metal ions, a similar pattern emerged as for CP-Ser and CP-Ser\(_\alpha\) at sub-stoichiometric levels of metal ions. At 0.1 equiv of Zn(II) or Cu(II), the ThT assay indicated that both S100B and S100B\(_\alpha\) had a small, but measurable influence on the amount of \(\beta\)-sheet-rich A\(\beta\)\(_{40}\) aggregates formed (Fig. 1b). On the other hand, the ThT assay with 1 equiv of Zn(II) or Cu(II) indicated that neither S100B nor S100B\(_\alpha\) could significantly alter the production of \(\beta\)-sheet-rich A\(\beta\)\(_{40}\) aggregates (Fig. 1b). These results were supported by TEM at 1 equiv of Zn(II) or Cu(II), which presented relatively small-sized fibrils, instead of larger and longer fibrils, that were distinct from S100B and S100B\(_\alpha\) aggregates (Fig. 2b). Although S100B binds transition metal ions, including Cu(II) and Zn(II), \(^{45,47,48,73}\) the protein may not be able to tune the aggregation of metal–A\(\beta\)\(_{40}\). Similar to CP-Ser and CP-Ser\(_\alpha\), neither S100B nor S100B\(_\alpha\) could interfere with fibrillation of metal-free A\(\beta\)\(_{42}\) and metal–A\(\beta\)\(_{42}\) (Fig. 4b and d, ESI\(^\dagger\)). Overall, S100B and S100B\(_\alpha\) were not observed to affect metal-free and metal-bound A\(\beta\)\(_{40}\) aggregation like CP.

Size distributions of metal-free A\(\beta\)\(_{40}\) and metal–A\(\beta\)\(_{40}\) aggregates generated upon treatment with S100 proteins

To assess the impact of these four S100 proteins on A\(\beta\)\(_{40}\) oligomer formation, gel/Western blot and dot blot analysis were performed to visualize the molecular weight (MW) distribution of A\(\beta\)\(_{40}\) species upon incubation of A\(\beta\)\(_{40}\) with each S100 protein (Fig. 3a and b and Fig. S6 and S7, ESI\(^\dagger\)). An anti-A\(\beta\)\(_{40}\) antibody (6E10) \(^{44,75}\) was employed to visualize A\(\beta\)\(_{40}\), an anti-S100A9 antibody was used to detect CP-Ser and CP-Ser\(_\alpha\), and Coomassie blue staining was employed to indicate S100B and S100B\(_\alpha\). Gel electrophoresis was conducted in running buffer containing 1% SDS to mobilize protein samples through the gel matrix. This amount of SDS likely disrupts complexes of S100 proteins with A\(\beta\)\(_{40}\) that may exist, but does not break-up SDS-resistant A\(\beta\)\(_{40}\) oligomers. Therefore, the bands observed in the 6E10 blots can be ascribed solely to A\(\beta\)\(_{40}\) oligomers (Fig. 3a...
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and Fig. S6 and S7, ESI†; gray), and this assignment is further supported by the anti-S100A9 blots and Coomassie blue stains since these data present almost identical patterns of S100 proteins in the absence and presence of Aβ (Fig. 3a and Fig. S6 and S7, ESI†). These data presented similar patterns of S100 proteins to those presented in our previous study. These results indicated that CP-Ser is not likely to be involved in S100 protein aggregation, and that CP-Ser might have an additional effect on Aβ aggregation.

Fig. 3 Analyses of the resultant metal-free Aβ40 species upon treatment with S100 proteins (CP-Ser, CP-Ser, S100B, and S100B). (a) Aβ40 species produced after 24 h incubation with or without the proteins, visualized by gel/Western blot using an anti-Aβ antibody (6E10; gray gels) or an anti-S100A9 antibody (blue gels; CP-Ser and CP-Ser) as well as Coomassie blue staining (blue gels; S100B and S100B). (b) Dot blot analysis of metal-free Aβ40 species generated after 24 h incubation with or without the proteins, monitored employing an anti-Aβ antibody (6E10), an anti-amyloid oligomer antibody (A11), and an anti-amyloid fibril antibody (OC). Samples: [Aβ40] = 20 µM; [S100 proteins] = 20 µM; pH 7.4; 24 h incubation; 37 °C.

S6 and S7 ESI†; blue). Clear bands representing monomeric S100A9 (ca. 13 kDa) and S100B (ca. 11 kDa) as well as the CP-Ser heterodimer (ca. 24 kDa) and the S100B homodimer (ca. 21 kDa) were observed.

Without metal ions, addition of CP-Ser to Aβ40 resulted in smaller oligomers compared to the oligomers in the samples containing Aβ40 only (Fig. 3a). This observation is in general agreement with the observations from the ThT assays and TEM (Fig. 1b and 2a), which together indicates that CP-Ser modulates the aggregation pathway of Aβ40. To obtain further insight into this process, the Aβ40 oligomers and fibrils formed upon incubation with CP-Ser were detected by dot blots using the amyloid-oligomer-targeting antibody (A11) and a fibril-reactive antibody (OC) (Fig. 3b). The intensity of the dots blotted with A11 were similar between samples of CP-Ser alone and CP-Ser with Aβ40. In contrast, both CP-Ser alone and CP-Ser with Aβ40 had low reactivity against OC (presented lower intensity than Aβ40 only). These results suggested that CP-Ser formed structured amyloid oligomers, but not fibrils, under our experimental conditions. Fibril formation of Aβ40 begins with a nucleation phase in which soluble Aβ40 monomers produce small oligomers that seed the generation of protofibrils in the elongation phase.42 CP-Ser, either as a heterodimer or higher-order oligomer, might interfere with the ability of Aβ40 oligomers to form a nucleus for fibrilization, potentially by forming complexes with Aβ40. Indeed, the anti-S100A9 blots (Fig. 3a) show a slight streaking pattern in the presence of 1% SDS, which may be evidence for formation of an Aβ40–CP-Ser complex.

CP-Ser also impacted the MW distribution of Aβ40 species (Fig. 3a and b). Dot blots showed that CP-Ser also responded to the A11 antibody, but not to the OC antibody (Fig. 3b), indicating no significance of the metal-binding sites in the formation of CP-Ser oligomers. Nevertheless, CP-Ser appeared to oligomerize or bind to Aβ40 less than CP-Ser as evidenced by the absence of streaking in the anti-S100A9 blot (Fig. 3a).

Both S100B and S100B exhibited a relatively subtle effect on the formation of Aβ40 oligomers. In the gel/Western blot with 6E10, both S100B and S100B exhibited smearing bands, compared to the samples of Aβ40 only (Fig. 3a). This impact on the generation of Aβ40 oligomers was also confirmed by the dot blot where A11-treated dots were more intense in the presence of S100B and S100B, than in the absence of the protein, especially for S100B. Moreover, the intensities of the dots blotted with the OC antibody for the samples of Aβ40 and S100B or S100B were similar to the sample of Aβ40 only. This observation is in agreement with both the ThT (Fig. 1b) and TEM analyses (Fig. 2b) presenting only a slightly discernible influence on Aβ fibril formation, which may not be resolvable by gel/Western and dot blots. The samples containing only S100B and S100B, did not react to the A11 antibody, which could indicate that structured amyloid oligomers were not produced upon incubation with these proteins under our experimental conditions and may be one contributing factor for the distinct behavior of CP-Ser and CP-Ser versus S100B and S100B, towards Aβ40 aggregation.

The size distribution of Aβ40 and S100 protein aggregates with different amounts of metal ions (i.e., 0.1 and 1.0 equiv of Zn(II) and Cu(II)) upon incubation was also investigated by gel/Western blot (Fig. S6 and S7, ESI†). The results generally agree with the data collected employing other methods (Fig. 1b and 2). One exception was the lack of change in Aβ40 oligomer distribution, observed by gel/Western blot, when CP-Ser was added to the samples containing Zn(II) (0.1 equiv). This observation contradicted the results from ThT and TEM that showed an influence of CP-Ser on Aβ fibril formation at sub-stoichiometric amounts of metal ions (Fig. 1b). This apparent discrepancy may be explained by the production of larger Aβ40 aggregates in the presence of CP-Ser, which are incapable of passing into the gel matrix and may be observed as a high intensity at the top of the gel within the well. Alternatively, the 1% SDS may dissociate the oligomeric forms of CP-Ser in the presence of metal ions under our experimental conditions (Fig. S6 and S7, ESI†). Another possible explanation could be that CP-Ser oligomers influence the population of metal-free Aβ40 when sub-stoichiometric metal ions were added, but might leave metal–Aβ40 species unaffected, which appeared in the gel as metal–Aβ40 samples without CP-Ser.

With S100B or S100B, at 1 equiv of Zn(II) or Cu(II) to Aβ40, new bands were observed by gel/Western blot (Fig. S6, ESI†). In the presence of Cu(II), S100B and S100B exhibited multiple
bands which correspond to the MW of the S100B homodimer (Fig. S6b and S7b, ESI†). S100B contains Cys residues, and these dimeric species may be generated by disulfide bond formation between S100B monomers, as a previous study suggested. These bands may also indicate that both S100B monomeric and dimeric Aβ can interact with Aβ40 and metal ions; however, these proteins may not have a significant effect on the aggregation or fibrilization of Aβ40 as indicated by the ThT assay (Fig. 1b) and TEM (Fig. 2b).

Interaction between CP-Ser or S100B and Aβ40 identified by ESI-MS

Because CP-Ser exhibited a pronounced effect on Aβ40 aggregation, we undertook further studies to determine whether a direct interaction between CP-Ser and Aβ40 could be observed in the form of complexes by ESI-MS. Mass spectra of the samples containing CP-Ser only and mixtures of CP-Ser and Aβ40 revealed the expected peaks attributable to the CP-Ser heterodimer (green) as well as monomers of S100A8 (yellow circles) and S100A9 (blue circles) (Fig. 4). The spectra obtained for CP-Ser and Aβ40 also contained peaks assigned to monomeric and dimeric Aβ40 (black circles and triangles, respectively), as well as peaks assigned to Aβ40–CP-Ser complexes (red circles) (Fig. 4). In addition to complex formation, the charge distributions of the CP-Ser peaks were altered in the presence of Aβ40, which may be triggered by the interactions between the two proteins (Fig. 4). Moreover, in the case of S100B, complex formation between S100B and Aβ40 was not observed under our experimental conditions (Fig. S8, ESI†), supporting a weaker interaction between the two proteins, as observed by the ThT assay (Fig. 1b) and TEM experiment (Fig. 2b). Overall, the direct interaction between CP-Ser and Aβ40 was observed, suggesting a connection between such protein-protein interaction and the ability of CP-Ser to generate less β-sheet-rich aggregates of Aβ40.

Cytotoxicity associated with CP-Ser in the absence and presence of Aβ40 and metal ions

Although cells secrete CP to protect themselves from pathogen invasion, elevated and uncontrolled levels of CP may have detrimental effects. To assess this possibility in the context of the current work, the viability of human neuroblastoma SH-SY5Y (SY) cells treated with CP-Ser was examined (Fig. 5a). CP-Ser was pre-incubated for 8 h in either the absence or presence of Aβ and applied to SY cells. After a 6 h incubation with SY cells, the CP-Ser species, pre-incubated in the absence of Aβ, did not cause noticeable toxicity, whereas a 24 h incubation afforded some toxicity (ca. 20% cell death) (Fig. 5a). Dot blots with the A11 antibody indicated that, in the presence of SY cells, CP-Ser formed structured oligomers at the 6 h time point, and that more A11-detectable species were observed at the longer, 24 h time point (Fig. 5b). In contrast, structured oligomers were not detected with the A11 antibody when the protein was incubated alone without cells in buffered solution for 8 h (Fig. S9, ESI†), indicating either that the CP-Ser did not aggregate or that it aggregated into amorphous aggregates unreactive to A11 antibody. These structured oligomeric species of CP-Ser may cause the observed toxicity.

CP-Ser combined with Aβ40 could generate slightly more toxic species than CP-Ser or Aβ40 alone (Fig. 5a). For instance, ca. 69% cell survival was observed after 6 h incubation with CP-Ser and Aβ40 compared to 96% and 83% cell survival for cell treated with CP-Ser or Aβ40 alone. These trends were also observed after 24 h incubation (Fig. 5a). When combined together in cell culture, CP-Ser and Aβ40 formed structured oligomeric species, detected by A11 (Fig. 5b), and these A11-reactive aggregates may exhibit cytotoxicity (Fig. 5a).

In the case of Aβ42, similar cytotoxicity was observed for the pre-incubated CP-Ser and Aβ42 mixture and Aβ42 alone after a 6 h incubation (ca. 77% cell viability; Fig. 5a). In the presence of added Zn(II) and Cu(II), greater than ca. 96% of cells survived upon 6 h incubation with pre-incubated (for 8 h) CP-Ser (Fig. S10, ESI†). Moreover, Zn(II)-treated CP-Ser showed no toxicity at the 24 h time point (ca. 100% cell survival), whereas CP-Ser with Cu(II) showed some toxicity (ca. 86% cell viability) (Fig. S10, ESI†). The cell viability of CP-Ser with metal ions and Aβ was relatively similar to that from the cells treated with metal ions and Aβ. Therefore, CP-Ser may not be able to affect the toxicity induced by metal-associated Aβ species significantly.
Overall, CP-Ser species may be toxic agents to cells exposed to the proteins longer than 24 h, either via the formation of Aβ-reactive structured oligomers or because they may sequester metal ions from the media since exogenous Zn(II) or Cu(II) mitigates toxicity. Additionally, the toxicity of CP-Ser species at 24 h could be slightly increased by interacting with Aβ40 in the absence of metal ions. Based on these cell studies, CP-Ser may not have significant influence on the toxicity induced by Aβ40 or Aβ42, with and without metal ions. Further studies are required to understand the toxicity caused by CP-Ser and CP-Ser–Aβ.

Conclusions

In this work, we examined the ability of CP-Ser and S100B to alter both metal-free Aβ and metal–Aβ aggregation. Out of the proteins examined, the influence of CP-Ser on Aβ40 aggregation was most pronounced. Such an impact on Aβ40 aggregation in the absence and presence of metal ions could potentially occur by the generation of complexes between CP-Ser and Aβ40, as observed by ESI-MS, or by the formation of CP-Ser oligomers that might alter the aggregation of Aβ40 and enhance the toxicity triggered by Aβ40.

The molecular details of how CP-Ser inhibits metal-free Aβ40 and metal–Aβ40 fibril formation are unknown. One possibility suggested by this work is that CP-Ser interacts directly with metal-free Aβ40 to alter Aβ40 aggregation, possibly via direct protein–protein interactions as supported by ESI-MS. Another potential mechanism is that CP-Ser influences metal–Aβ40 aggregation by disrupting metal binding to Aβ40, which is supported by the observation that CP-Ser(Aβ, which lacks defined metal binding sites, indicated no influence on Aβ aggregation when ≥ 1 equiv of Zn(II) and Cu(II) were added under the assay conditions. Therefore, the impact of CP-Ser could be dual in nature, by directly interacting with Aβ40 to limit aggregation and by interfering metal binding to Aβ40 to attenuate metal-induced Aβ40 aggregation, which was revealed by comparing the behavior of CP-Ser to CP-Ser(Aβ.

By demonstrating the interplay between CP-Ser and Aβ, our studies suggest that CP and Aβ may interact in vivo. If indeed the case, this work also supports the increasingly appreciated multi-faceted nature of AD pathology, wherein several factors may be involved in the progression of the disease.

In closing, protein–protein networks should be identified to gain a better understanding of AD pathogenesis.

Experimental

Materials and methods

All reagents were purchased from commercial suppliers and used as received unless otherwise stated. CP-Ser and S100B proteins were overexpressed, purified, and characterized as previously reported. Trace metal contamination was removed from buffers and solutions used for Aβ experiments by treating with Chelex (Sigma-Aldrich, St. Louis, MO, USA). Aβ40 (DAEFRHDSGYEVHHQKLVFFAEDVGSNKGAIIGLMVGGVV), and Aβ42 (DAEFRHDSGYEVHHQKLVFFAEDVGSNKGAIIGLMVGGVV) were purchased from Anaspec (Fremont, CA, USA). We have carried out the experiments with more than three replicates with at least two different batches of Aβ. All double distilled H2O (ddH2O) used during experiments was obtained from a Milli-Q Direct 16 system (Merck KGaA, Darmstadt, Germany). TEM images were recorded on a Phillips CM-100 transmission electron microscope [Microscopy and Image Analysis Laboratory (MIL), University of Michigan, Ann Arbor, MI, USA] or a JEOL JEM-2100 transmission electron microscope [UNIST Central Research Facilities (UCRF), Ulsan, Republic of Korea]. A SpectraMax M5 microplate reader (Molecular Devices, Sunnyvale, CA, USA) was used to measure the fluorescence intensity and absorbance for the assays.

Protein preparation and purification

The S100A8 and S100A9 subunits for CP-Ser and CP-Ser(Aβ were overexpressed in Escherichia coli BL21 (DE3) and the proteins were reconstituted, purified, and stored at −80 °C according to previously established methods. The procedure gives the apo heterodimers. Each protein sample was freeze-thawed only once before use. S100B was overexpressed in Escherichia coli BL21 (DE3) and purified as previously described. Procedures

![Fig. 5](image_url)

Fig. 5 Determination of the toxicity of CP-Ser with and without Aβ in cells and their conformational state by dot blot. (a) Viability of 5Y cells incubated with CP-Ser and/or Aβ. CP-Ser was pre-incubated in both the absence and presence of Aβ for 8 h and incubated with 5Y cells for 6 (left) or 24 h (right). Cytotoxicity was measured by the MTT assay. The viability values were calculated by comparison to cells treated with a volume of H2O equal to the protein added. Error bars represent the standard error from three independent experiments. Conditions (final concentrations): [CP-Ser] = 5 μM; [Aβ] = 5 μM. (b) Dot blot assay of pre-incubated (for 8 h) CP-Ser with and without Aβ after 6 (left) or 24 h (right) incubation with cells. Conditions: [CP-Ser] = 5 μM; [Aβ] = 5 μM.
for the expression and purification of S100B, are provided as Supporting Information.

**Aβ aggregation experiments**

All experiments with Aβ were performed according to previously published methods. Aβ peptides were dissolved in ammonium hydroxide (NH₄OH, 1% v/v,aq), aliquoted, lyophilized, and stored at −80 °C. A stock solution (ca. 200 µM) was prepared by dissolving Aβ in NH₄OH (1% w/v, aq, 10 µl) followed by dilution with ddH₂O, as reported previously. The Aβ₄₀ concentration was determined by measuring the absorbance of the solution at 280 nm (ε = 1450 M⁻¹ cm⁻¹ for Aβ₄₀). HEPES [4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid; 20 mM, pH 7.4 (for metal-free and Zn(II) samples) and pH 6.6 (for Cu(II) samples), 150 mM NaCl was used for the studies. For the experiments, freshly dissolved Aβ (20 µM) in the absence and presence of a metal chloride salt (ZnCl₂) was treated with S100 proteins (CP-Ser, CP-Ser₃₄, S100B, and S100B₃) at 20 µM and incubated for 24 h at 37 °C with constant agitation.

**ThT assay**

The amount of Aβ fibrillization and the kinetics of Aβ₄₀ aggregation, treated with S100 proteins (20 µM) with or without metal ions (ZnCl₂ or CuCl₂; 2 or 20 µM) and S100 proteins (CP-Ser, CP-Ser₃₄, S100B, and S100B₃; 20 µM) was obtained after 24 h incubation at 37 °C with constant agitation and diluted with buffer (20 mM HEPES, pH 7.4 (for metal-free and Zn(II) experiments) or pH 6.6 (for Cu(II) experiments), 150 mM NaCl). For monitoring Aβ₄₀ aggregation kinetics, each Aβ sample (20 µM) with or without CP-Ser was obtained after 0, 0.3, 0.6, 1, 2, 4, 8, 12, and 24 h incubation at 37 °C with constant agitation. The samples were incubated for 20 min with ThT (20 µM) and the fluorescence intensity was measured by a microplate reader (λₑm = 440 nm; λₑx = 490 nm) and normalized relative to that of metal-free Aβ₄₀ aggregates (without CP-Ser) generated by 24 h incubation. The ThT signal was determined at least 20 min at room temperature. Images of samples were discharged grids (Formvar/Carbon 300-mesh, Electron Microscopy Sciences, Hatfield, PA, USA) were treated with samples from the experiments (5 µL) for 2 min at room temperature. Excess sample was removed with filter paper and the grids were washed with ddH₂O three times. Each grid was stained with uranyl acetate (1% w/v ddH₂O; 5 µL) for 1 min. Uranyl acetate was blotted on and grids were dried for at least 20 min at room temperature. Images of samples were obtained after 24 h incubation at 37 °C with constant agitation.

**Gel electrophoresis with Western blot**

The Aβ peptide experiments described above were analyzed by gel electrophoresis with two different visualization methods: (i) Western blotting (gel/Western blot) with two different antibodies (anti-Aβ antibody (6E10; Covance, Princeton, NJ, USA) or anti-S100A9 antibody (Santa Cruz Biotechnology, Dallas, TX, USA));²³,⁷⁵,⁸²–⁸⁴ (ii) Coomassie blue staining. Each sample from the experiments was separated using a 10–20% gradient Tris-tricine gel (Thermo Fisher, Grand Island, NY, USA). First, the CP-Ser samples were separated on the gel, transferred to a nitrocellulose membrane, and blocked for 3 h at room temperature with a bovine serum albumin (BSA) solution (3% w/v) in Tris-buffered saline (TBS, Fisher, Pittsburgh, PA, USA) containing 0.1% Tween-20 (Sigma-Aldrich; TBS-T). The membrane was treated with the Aβ monoclonal antibody (6E10, 1:2,000, BSA 2% w/v, in TBS-T) or anti-S100A9 antibody (1:1000, BSA 2% w/v, in TBS-T) overnight at 4 °C and then probed with a horseradish peroxidase conjugated goat anti-mouse secondary antibody (1:5,000; Cappel Chemical, Ann Arbor, MI, USA) or donkey anti-goat secondary antibody (1:5,000; Jackson ImmunoResearch Laboratories, West Grove, PA, USA) in 2% w/v BSA (in TBS-T) solution for 1 h at room temperature against anti-Aβ and anti-S100A9, respectively. The protein bands were visualized using Thermo Scientific Supersignal West Pico Chemiluminescent Substrate (Rockford, IL, USA) or a homemade ECL kit²⁸ was used to visualize the results on a ChemiDoc MP Imaging System (Bio-rad, Hercules, CA, USA).

The gels run with S100B- and S100B₃-containing samples were stained with Coomassie blue for 30 min and destained for 3 h. Gel images were obtained by a ChemiDoc MP system (Bio-rad). Then, the dye was completely destained from the gel, and the gel was transferred to a nitrocellulose membrane for Western blot with 6E10 (1:2,000) in a solution of 2% w/v BSA (in TBS-T) and a horseradish peroxidase conjugated goat anti-mouse secondary antibody (1:5,000) same as the Western blot for the CP-Ser samples.

**Dot blot assay**

Solutions of protein aggregates (2 µL) from the experiments were spotted on a nitrocellulose membrane and the membrane was blocked with BSA solution (3% w/v; RMBIO, Missoula, MT, USA) in Tris-buffered saline containing 0.01% Tween 20 (TBS-T) at room temperature for 1.5 h. Then, the membrane was incubated with a primary antibody, 6E10 (1:2,000), A11 (1:2,500; Millipore, Billerica, MA, USA) or OC (1:2,500; Millipore) in a solution of 2% w/v BSA (in TBS-T) for 1.5 h at room temperature. After washing with TBS-T three times (7 min each), the horseradish peroxidase-conjugated goat anti-mouse (for 6E10) or goat anti-rabbit (for A11 and OC) secondary antibody (1:5,000; Cappel Chemical Company for 6E10, 1:2,500; Promega for A11 and OC) in the solution of BSA (2% w/v in TBS-T) was added to the membrane and incubated for 1 h at room temperature. A homemade ECL kit²₈ was used to visualize the results on a ChemiDoc MP Imaging System (Bio-rad). The same membrane was stripped by treating with hydrogen peroxide (H₂O₂) for 30 min at room temperature, washed 4 times with TBS-T for 10 min each, blocked with the solution of BSA (3% w/v in TBS-T), and incubated with the primary antibodies (6E10, A11, and OC) as described above.

**Transmission electron microscopy (TEM)**

Protein samples for TEM measurements were prepared following previously reported methods.³⁴ Glow discharged grids (Formvar/Carbon 300-mesh, Electron Microscopy Sciences, Hatfield, PA, USA) were treated with samples from the experiments (5 µL) for 2 min at room temperature. Excess sample was removed with filter paper and the grids were washed with ddH₂O three times. Each grid was stained with uranyl acetate (1% w/v ddH₂O; 5 µL) for 1 min. Uranyl acetate was blotted on and grids were dried for at least 20 min at room temperature. Images of samples were obtained after 24 h incubation at 37 °C with constant agitation.
taken by a Philips CM-100 transmission electron microscope (80 kV, 25,000x magnification; MIL, University of Michigan, Ann Arbor, MI, USA) or JEOL JEM-2100 transmission electron microscope (200 kV, 25,000x magnification; UCRF, Ulsan, Republic of Korea).

Mass spectrometric studies

Mass spectrometric experiments with CP-Ser and Aβ40 were performed on a Synapt G2-Si Q-ToF mass spectrometer equipped with electrospray ionization (ESI) source (Waters, Manchester, UK). Aβ40 (20 µM) was mixed with CP-Ser (20 µM) or S100B (20 µM) in 10 mM ammonium acetate (pH 7.5). The prepared samples were incubated at 37 °C for 24 h without agitation. Incubated samples were diluted 4-fold with 10 mM ammonium acetate (pH 7.5) before injection, and the final concentration of peptides was adjusted to 5 µM. The capillary voltage, sampling cone voltage, and source temperature were adjusted to 2.8 kV, 70 V, and 40 °C, respectively, for CP-Ser and 1.8 kV, 40 V, and 40 °C, respectively, for S100B. Moreover, for the competition experiments, the samples were prepared in 10 mM ammonium acetate (pH 7.5) through two different addition orders of Aβ40, CuCl2, and CP-Ser: (i) Aβ40 (20 µM) was first treated with CuCl2 (20 µM) followed by addition of CP-Ser (20 µM); (ii) CP-Ser (20 µM) was first incubated with CuCl2 followed by treatment with Aβ40. The prepared samples were incubated at 37 °C for 1 h without agitation, and then were diluted by four fold with 10 mM ammonium acetate (pH 7.5) (the final concentration of peptides was adjusted to 5 µM) before injection to the mass spectrometer. The capillary voltage, sampling cone voltage, and source temperature were adjusted to 1.8 kV, 40 V, and 40 °C, respectively. As the resolution of the instrument was not high enough to determine the monoisotopic masses for CP-Ser subunits, heterodimeric CP-Ser, or homodimeric S100B complexes with Aβ40, all peaks were assigned using the m/z values of the most abundant isotopes. For example, if the m/z of an apex isotope is divided by the mass of the proteins and the result is an integer, the peak can be assigned as one charge state of the proteins.

Cell viability measurements

The SH-SY5Y (SY) cell line was purchased from the American Type Culture Collection (ATCC, Manassas, VA, USA). The cell line was maintained in media containing 50% minimum essential medium (MEM) and 50% F12 (GIBCO), and supplemented with 10% fetal bovine serum (Sigma-Aldrich), and 100 U/mL penicillin (GIBCO). Cells were grown and maintained at 37 °C in a humidified atmosphere with 5% CO2. The cell culture used in this work did not indicate mycoplasma contamination. Cell viability upon treatment with CP-Ser was determined by the MTT assay [MTT = 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide]. Cells were seeded in a 96 well plate (13,000 cells in 100 µL per well). Cells were treated with pre-incubated proteins for 8 h with and without metal ions at 37 °C with constant agitation. After 6 or 24 h incubation with cells, MTT (25 µL of 5 mg/mL in PBS (pH 7.4, GIBCO)) was added to each well, and the plate was incubated for 4 h at 37 °C. Formazan produced by cells was solubilized using an acidic solution of N,N-dimethylformamide (DMF, pH 4.5, 50% v/v, aq) and sodium dodecyl sulfate (SDS, 20% w/v) overnight at room temperature in the dark. The absorbance was measured at 600 nm by a microplate reader. Cell viability was calculated relative to cells containing a volume of H2O equal to the volume of protein added.

Conflicts of interest

There are no conflicts to declare.

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