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DNA intercalators as amyloid assembly modulators: mechanistic insights†

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Amyloid assemblies are stabilized through inter-molecular H-bonding with initial structural organizations governed by extensive hydrophobic and π -stacking interactions. Apparently, the structural integrity and stability of DNA duplexes are also governed by a cooperative balance of similar interactions. Structural perturbations in both amyloids and DNA by planar molecules or intercalators essentially rely on their ability to interfere with this balance. However, poor information on amyloid interference mechanisms by intercalators has subjected researchers to undertake random trials with such anti-amyloidogenic agents. Here, we employed four different classes of DNA intercalators to investigate if their non-native, hetero-aromatic associations could modulate the amyloid aggregation pathway. We utilized micro-second scale MD simulations using the steric-zipper structure of the diabetes associated amylin fragment to identify such associations and their plausible role in subjugating higher order assemblies. The simulation estimates were experimentally validated and extended to other disease-associated amyloid systems including gelsolin, prion and lysozyme. We find that the intercalators essentially stabilize monomeric and prefibrillar assemblies, reducing their ability to transform into structured supramolecular structures. Our results conclusively establish the dominant role of aromatic-associations in diverting the course of the amyloid assembly process at the expense of stabilizing H-bonding networks. Overall this report presents comprehensive experimental and theoretical insights on the regulation of amyloidogenesis by aromatic planar moieties.

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Introduction

Amyloids represent fibrillar, intractable protein aggregates, the deposition of which in tissues causes a multitude of pathological complications including fatal neurodegenerative disorders such as Alzheimer's, Huntington's and Creutzfeldt Jakob diseases.^{1–4} Regardless of origin, sequence or native structure, amyloid aggregates share common cross- β spine architectures, primarily stabilized through inter-molecular hydrogen bond networks.^{5,6} Amyloidogenesis involves the destabilization of native protein architecture to form partially unfolded ensembles that attain rogue β -rich, oligomeric protofibrillar conformations.^{7,8} Self-association of protofibrillar intermediates terminates in ordered mature amyloid fibrils.^{9,10} Protofibrillar oligomer intermediates represent the central players in amyloidogenesis and amyloidosis. Structurally, they comprise fundamental elements for fibril elongation ('on pathway intermediates')^{11,12} and pathologically, account for amyloid

related cytotoxicities owing to their interaction with cellular membranes.^{13,14}

Transition of native monomers to protofibrillar intermediates depends on specific free surfaces formed by aggregation prone regions of proteins.^{15–17} These surfaces are defined through the intrinsic hydrophobicity, secondary structural propensities, and electrostatic nature of the constituent residues. The transitions along the aggregation pathway stem from complex interplay among these interactions in driving the assembly process. Amongst these interactions, recent literature on *in vitro* as well as *in silico* fibrillation of amyloidogenic regions from human islet amyloid polypeptide (IAP) projects synergistic role of π -stacking interactions, mediated through aromatic Phe's in the assembly process.^{18–21}

Similarly from the structural perspective, integrity and organization of DNA polymer rely on subtle balance of similar stabilizing H-bond and π -stacking interactions.^{22,23} Consequently, various planar molecules/compounds/drugs which are known to interact and modulate DNA structure have been tested for their anti-amyloidogenic potential.^{24–26} The best known drugs of this class included two FDA approved DNA intercalators, 4'-iodo-4'-deoxydoxorubicin for systemic amyloidosis^{27,28} and mitoxantrone/picoxantrone against A β (1–42) fibrillation.^{29,30} Traditionally, these compounds comprise

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simple class of planar organic molecules capable of modulating genetic and epigenetic expression *via* stacking in between adjacent base pairs of DNA.^{31–33} These interactions are specifically driven by pi stacking between the electron clouds of coplanar bases and the intercalators.^{34,35} Primarily hydrophobic interactions and beta sheet propensities have been proposed as main driving force during amyloid assembly.^{5,16} However, the finding that DNA intercalators like doxorubicin/mitoxantrone act as amyloid inhibitor warrants investigation of the balance between hydrophobic/beta-sheet propensities *vs.* aromatic interactions in driving amyloid assembly. In other words, whether the anti-amyloidogenic effect of the intercalators is due to their ability to favour non-amyloid ('off-pathway') intermediates through competitive "aromatic-associations" is yet to be proven.^{21,25}

Bridging the previously reported evidences, we hypothesize dominant role of hetero-aromatic associations in modulating amyloid aggregation pathways. During initial phases, such associations could interfere with inter-molecular interactions, obviating organization of native monomers to 'on-pathway' amyloid intermediates while driving the process towards 'off-pathway' intermediates. Additionally, such associations could also interfere with lateral assembly of protofibrillar intermediates, originally facilitated through π -associations, thereby impeding their progression to mature fibrils.^{36,37} Amyloid assembly in case of PolyQ is an exception as charge interactions appears to play dominant role. However, inhibition by EGCG by virtue of its hydrophobic/H-bond interactions requires thorough investigation behind its mechanism of action. To comprehensively understand the role of hetero-aromatic associations to interfere with the amyloidogenic pathways, we designed this current study by employing the pre-fibrillar structural coordinates of IAP amyloid-core containing sequence (NFGAILS), IAP (22–28) as our model system (Fig. 1). The choice of this system serves dual advantages in terms of Phe-23 mediated assembly formation^{18,38} and in pre-availability

of atomic coordinates of supramolecular amyloid assembly of IAP (22–28).³⁹ Thus, the entire reaction pathway for IAP (22–28) aggregation and in process interference mechanisms by planar compounds can be mapped through strategic simulation of its aggregation pathway.

For our studies we employed four different DNA intercalators: synthetic molecule, ethidium bromide (EtBr); natural alkaloid chelerythrine (Chl); and two FDA approved anti-cancer drugs *viz.*, doxorubicin (Dxr) and mitoxantrone (Mtx). Aggregation of IAP (22–28) was probed in the presence and absence of these intercalators by molecular dynamics simulations. Furthermore, the study was extended to investigate the effect of these compounds on the pre-formed protofibril assembly of IAP (22–28). To validate computational outcomes, this hetero-molecular landscape was replicated *in vitro* through spectroscopic approaches with visualization using electron microscopy. To eliminate selection bias towards IAP (22–28) model system, we employed three additional assorted systems, gelsolin peptide fragment (GNCFILD) from D187N disease mutant of gelsolin protein (AGel (186–192)) and other amyloidogenic proteins *viz.* lysozyme (Lys) and human prion protein (PrP). Gelsolin fragments and its mutations are associated with familial amyloidosis of Finnish type,⁴⁰ lysozyme in senile amyloidosis⁴¹ and prion aggregation in transmissible spongiform encephalopathies.⁴² Dissecting through diverse biophysical and visualization approaches on the same systems, our findings attempts to illustrate and correlate *in silico* landscapes with experimental estimates.

Results

Intercalators interrupt structural transitions into higher order assemblies

Molecular dynamic trajectories for each system were analyzed by distinct approaches. The amyloid modulatory function of intercalators was established by comparing the resultant

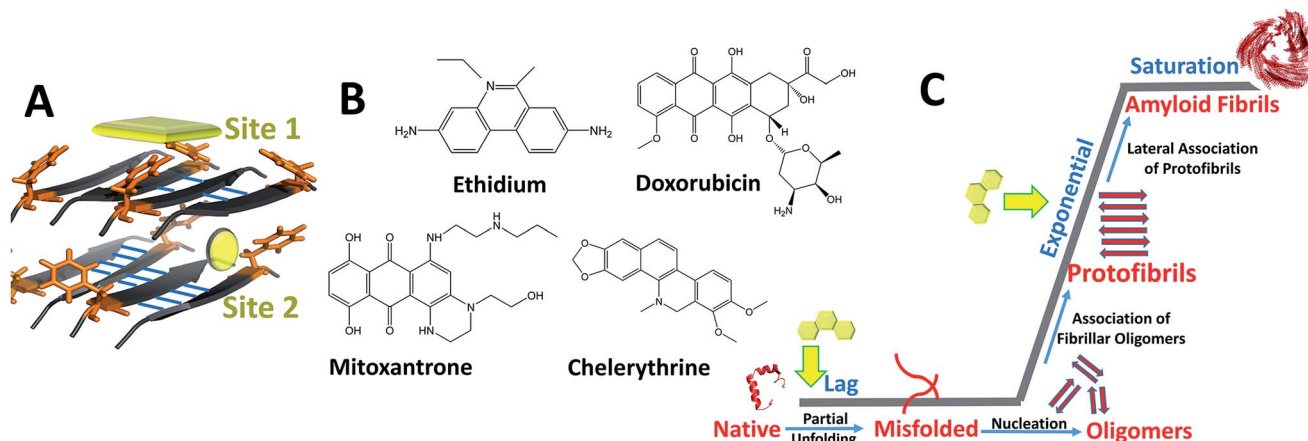


Fig. 1 Design of the present work (A) solid state NMR structure of IAP (22–28) protofibril assembly displaying aromatic Phe-23 residues (orange) at fibril surface (site 1) and within cross β -spine (site 2) and inter-molecular hydrogen pattern (blue). (B) chemical structures of DNA intercalators employed to study hetero-aromatic associations (C) generic aggregation pathway showing transitions of native monomer to amyloid fibrils through association of protofibrillar intermediates (dominant during exponential growth phase). Yellow arrows indicate target stages in this study to assess interference of intercalators with monomer to protofibril transitions and protofibril–protofibril associations to mature fibrils.



trajectories over six variables: root mean square deviation in distance between $C\alpha$ atoms of peptides (dRMS), hydrophobic solvent accessible surface area of side chains (hSAS), backbone secondary structure contents (SS), H-bond number (HBN), matrix of least distance between pair of residues (MLDR) and finally by free energy landscape (FEL) of aggregation pathways (Fig. 2 and 3). In our simulations the aggregates formed by IAP (22–28) and AGel (186–192) monomers in the absence of intercalators served as controls while the octameric protofibrillar assembly of IAP (22–28) and AGel (186–192) served as reference frames for respective systems (discussed in Materials and methodologies).

Aggregation landscape of IAP (22–28)

Variations in dRMS and hSAS. In the absence of compounds, transition of monomers to an oligomeric assembly was characterized by marked reduction in dRMS (Fig. 2A). Towards the end of simulation, the final frames approached gradually towards protofibril like oligomer assembly with ~ 0.2 Å dRMS from reference. Contrastingly, in the presence of compounds the dRMS remained persistently high (~ 0.8 Å), suggesting that the structures failed to converge into fibril like assembly. Transition of monomers to stable aggregates is accompanied by de-solvation of non-polar residues to form the amyloid core.^{43,44} Such transition can be monitored by changes in hSAS along the reaction coordinate.⁴⁴ In control, the observation of a steep drop in hSAS within 50 ns was indicative of early burial of monomer side chains through hydrophobic interactions followed by a gradual decline till the end of simulation (Fig. 2B). Presence of compounds resulted in comparatively lower drop (nearly 20 nm²) in hSAS compared to control suggesting substantially low percentage burial (Fig. 2B).

Variations in MLDR. In formation of IAP (22–28) assembly, formation of specific inter-residue contacts were represented as matrix of least distance between pair of residues (MLDR) averaged over whole trajectory (ESI Fig. 2†). The resultant matrix clearly indicated formation of large number of mutual native contacts between residues. In the presence of intercalators, the respective MLDR's clearly depicted augmentation in average distance among the residues clearly indicating integration of intercalators along the monomeric units, resulting in higher inter residue distance.

Variations in HBN and SS. Furthermore, reduction in inter-residue contacts guided towards the possibility of the intercalators interfering with inter-peptide H-bond network (HBN), the principal interactions in stabilizing amyloid aggregates. Presence of intercalators resulted in lower HBN (<33 for EtBr, Mtx, Chl and <30 for Dxr per time frame) as compared to control (>38 per time frame) averaged over last 400 ns of simulation (Fig. 2C).

The fully ordered octameric protofibril like assembly (2kib) possesses nearly 70% residues with high propensity for extended conformation.⁴⁵ In our study, we observed a progressive increase in the beta sheet content with a concomitant decline in coil content in the control samples ($\sim 35\%$ increase and $\sim 50\%$ decrease respectively, Fig. 2D). In the presence of intercalators, the aggregates displayed an overall higher random conformation for majority of residues, thus exhibiting a strong structural disorder. Apart from intrinsic hydrophobicity and β -sheet propensities, IAP (22–28) aggregation pathway had been shown to be driven by homo-aromatic associations among its Phe-23 residues.¹⁸ Therefore, we speculated subsistence of hetero-aromatic associations between Phe-23

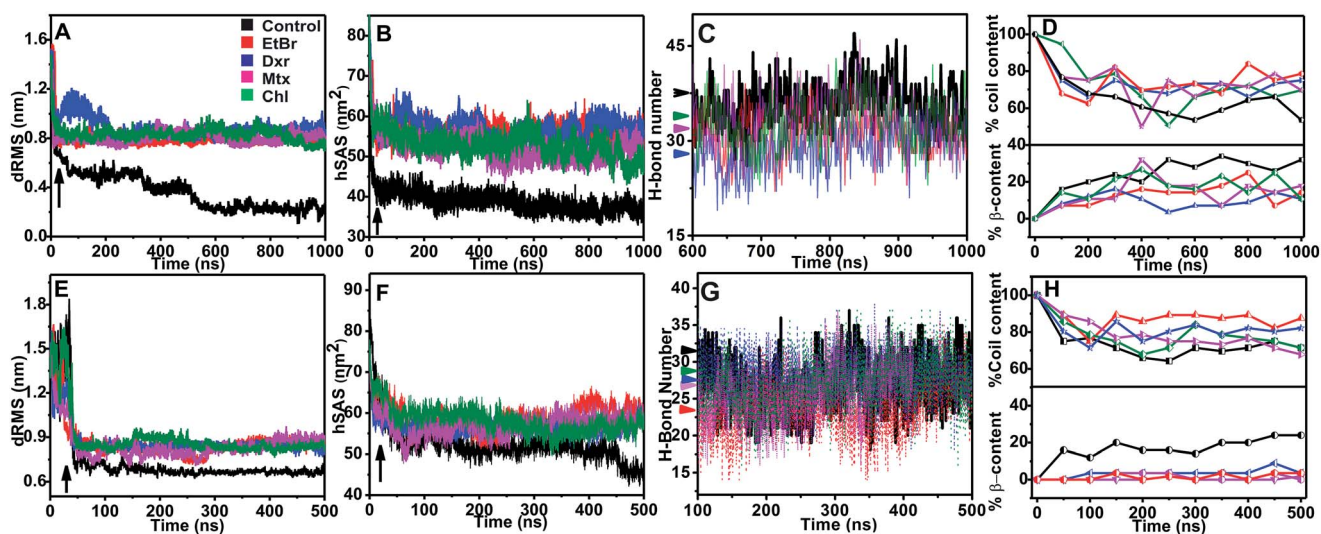


Fig. 2 Time development of variables for IAP (22–28) and AGel (186–192) aggregation. Variations in dRMS, hSAS, HBN and SS contents for transitions of IAP (22–28) (A–D) and AGel (186–192) (E–H) monomers to oligomeric protofibrillar assembly in the absence (black) and in the presence of different intercalators (red: EtBr, blue: Dxr, green: Mtx and Chl: magenta) during our simulation runs of 1000 ns and 500 ns respectively. Formation of oligomeric protofibrillar assemblies from peptide monomers can be deduced through marginal deviation of dRMS (A and E) and hSAS (B and F) from reference protofibrillar assemblies. In the presence of intercalators, dRMS and hSAS ceased to converge, indicating formation of non-amyloid like assemblies. This was confirmed through their reduced HBN (C and G) and SS contents (D and H) compared to control systems (black trajectories). Black arrows in A, B and E, F signifies event where hydrophobic collapse was formed by IAP (22–28) and AGel (186–192) monomers respectively. Respective colored arrows in C and G represents average number of H-bonds per time frame.



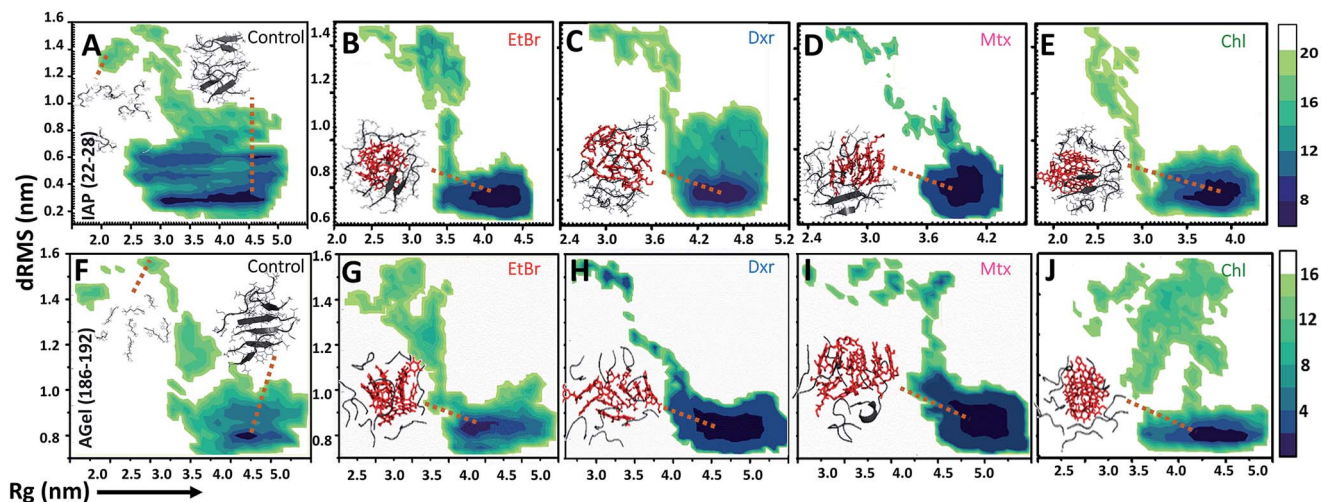


Fig. 3 Mapping aggregation pathway of IAP (22–28) and AGel (186–192). Free energy landscapes (in kcal mol⁻¹) of IAP (22–28) and AGel (186–192) aggregation in the absence (A and F respectively) and presence (B–E and G–J) of EtBr, Dxr, Mtx and Chl respectively. Conformations representing lowest free energy states (grey cartoons) both in their absence and presence are shown as insets (intercalators are shown as red involved in interaction with monomeric units).

and planar compounds driving the pathway towards non-amyloid assemblies. The MLDRs constructed between Phe-23 and test compounds for each system averaged over final 600 ns of trajectory showed persistent hetero-aromatic associations (ESI Fig. 3B–3E†). Converging the clues from H-bond patterns, β -content evolution and hetero-aromatic associations, we confirmed that intercalators had deviated aggregation of IAP (22–28) to non-amyloid assemblies, putatively incapable of organizing into higher order amyloid like assemblies.

Free energy landscapes. Finally, free energy landscapes (FEL) were constructed to sample preferred conformational space of any assembly formed in the absence and presence of four intercalators (Fig. 3), projected as function of gyration radius (R_g) and dRMS. The FEL of control IAP (22–28) aggregation clearly displayed formation of ensembles occupying most of the conformational space comprising broad, low energy basin. These could be characterized by ensembles undergoing intra molecular rearrangements to form stable assembly. However, the lowest energy conformations were restricted to narrow basin representing partially ordered compact oligomeric assembly (Fig. 3A). In contrast, presence of intercalators altered the aggregation landscape of IAP (22–28) with lowest free energy conformations, mostly confined to single large basin towards the boundary of the landscape. These conformations were characterized by disordered and less compact structures, inferred from their relatively higher average R_g (1.5–2) times, compared to the R_g of the assembly in control. The representative structures corresponding to lowest energy conformations of the non-amyloid assembly are shown as insets in Fig. 3B–E.

Aggregation landscape of AGel (186–192)

The influence of intercalators was further investigated on aggregation landscape of AGel (186–192). Here, moderate disparity in dRMS, hSAS and HBN was observed between the

control and the test samples (Fig. 2E–H). The average β -sheet content in control systems progressively escalated with 25% of residues involved in forming in-register beta arrangement. In the presence of intercalators, each system failed to organize into β -rich aggregates throughout the simulation (Fig. 2H). Respective MLDRs showed that fraction of native contacts were subsided by intercalators compared to control (ESI Fig. 4†). The employed intercalators similarly displayed hetero-aromatic associations with Phe residues of AGel (186–192) monomers along the reaction pathway (ESI Fig. 5†). The most stable conformations during aggregation pathways were projected as function of R_g and dRMS variations with respect to reference structure. The FEL of control system displayed spontaneous assembly of monomers, without the accumulation of intermediate states; to yield β -rich aggregates (Fig. 3F). The corresponding lowest energy conformations were confined to narrow basin, possessing higher R_g (3.1 nm) compared to the reference ($R_g \sim 2.5$ nm). A notable finding on extracting structures from the coordinates of basin was formation of predominantly single layer heterogeneous and in-register fibril assembly resulting in higher R_g than reference structure (double layer oligomer). Expectedly, the intercalators induced more disorder in the packing arrangement of monomers ($R_g \sim 4.5$ –5 nm, Fig. 3G–J) suggesting their integration within the monomeric units as seen in the IAP (22–28) simulations.

Intercalators remodel protofibrillar assembly

In order to assess if the intercalators could perturb the conformational integrity of pre-formed amyloid like ensembles, we carried out studies on protofibril like assembly of IAP (22–28). Initial docking studies through MOE showed two high propensity ligand binding sites (site 1 and 2, Fig. 1A) on assembly which showed comparable binding affinities (ESI Table 1†), hinting towards the probability of concurrent binding



at both sites. As a result, unbiased systems were setup to follow unrestrained protofibril-intercalator dynamics (ESI Fig. 1B†).

Variations in dRMS, hSAS and R_g

The effects of compounds on fibril assembly were determined in terms of comparative dRMS, hSAS and the overall structural changes in assembly (R_g) (Fig. 4). Control setup represented fibril assembly alone which showed progressive increase in dRMS, stabilizing near 0.45 Å post 400 ns simulation (Fig. 4A). This could be attributed to non-inclusion of dihedral constrains in these simulations, resulting in bending and twisting of protofibril assembly along its longitudinal axis. However, in the presence of EtBr and Dxr, the fibril assembly showed rapid fluctuations in dRMS around 550 ns and 850 ns respectively (Fig. 4A). The effect was more pronounced in the presence of EtBr where dRMS escalated to ~ 0.5 Å, comparably higher than control setup. Contrastingly, these fluctuations were completely absent in the presence of Mtx/Chl which displayed considerably lower dRMS (~ 0.3 Å) over entire trajectory indicating interactions primarily confined to fibril surface (site 1), thus restricting bending and twisting of fibrils. However, the interactions also resulted in slight unpacking of the assembly, indicated through R_g variations collected during last 500 ns of simulation period. Association of EtBr molecules with the assembly resulted in modulating fibril packing, indicated through comparatively high gyration radius (R_g) than control and in the presence of other intercalators (Fig. 4B). Furthermore, the much pronounced destabilization effects of EtBr on IAP (22–28) assembly were evident through concurrent solvation of

hydrophobic core residues (Fig. 4C). Dxr exerted mild solvation of the residues, only towards the end of simulation in accord with its dRMS fluctuations. Interestingly, Mtx and Chl facilitate sidechain packing of hydrophobic residues as observed by lower hSAS than control, contributing to higher percentage burial.

Hetero-aromatic associations and free-energy landscapes

In addition to H-bonding network, we found that hetero-aromatic associations between Phe residues of monomers and planar hydrophobic moiety of intercalators also played a crucial role. In control, unusual homo associations between Phe 7 and 8 and Phe 6 and 5 were observed (ESI Fig. 6†), possibly arising due to twisting and bending. Interestingly, EtBr completely hindered such homo aromatic associations and fostered hetero-aromatic associations through its phenanthridinium moiety. Each EtBr molecule displayed association not only with exposed Phe's (2, 4, 6 and 8) at fibril surface but also with Phe's (1, 3, 5 and 7) at site 2. Unlike EtBr, hetero-aromatic associations with other intercalators were less pronounced (ESI Fig. 6B†).

The influence of intercalators on the structural remodeling of pre-formed protofibrils into energetically favorable conformation was assessed by constructing FEL's. Fig. 4D shows projections of control assembly in the conformational space as function of R_g and backbone RMSD. The energetically favorable ensembles were confined to narrow conformational space, corresponding to compact (R_g 1.05 nm) twisted assembly. In the presence of EtBr and Dxr, the conformational space drifted towards partially condensed states (Fig. 4E and F). The preferred lowest energy conformations in the presence of EtBr

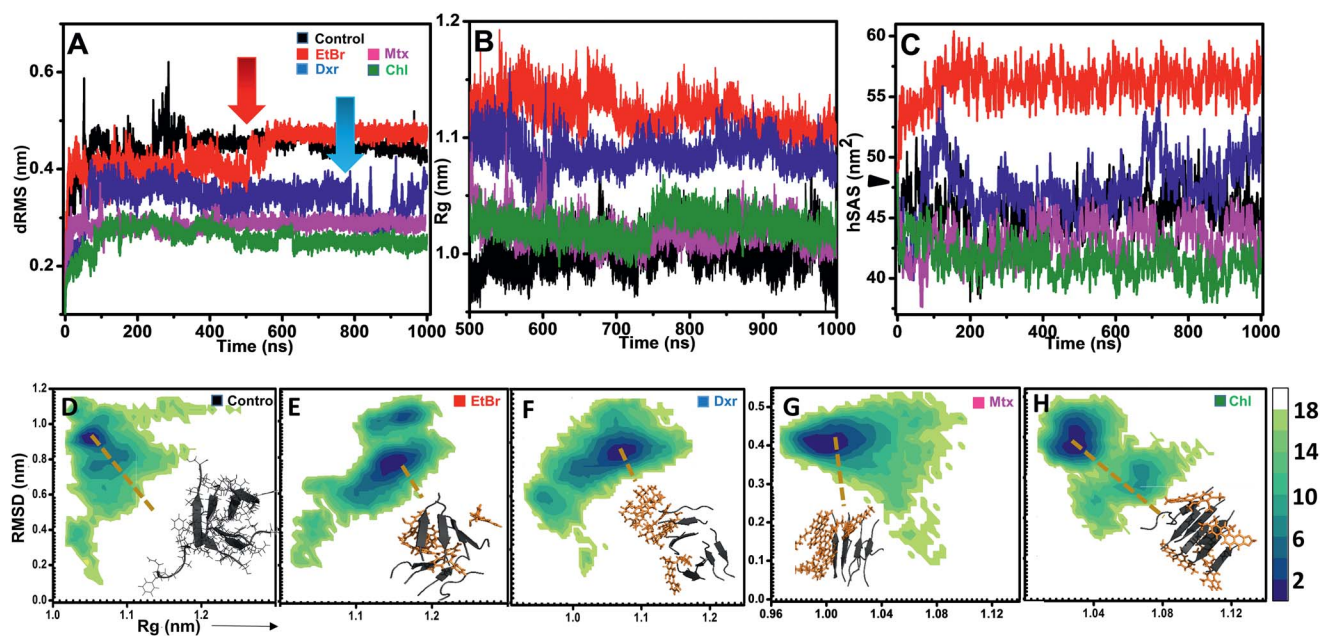


Fig. 4 Time development of variables during IAP (22–28) assembly modulation. Variations in dRMS (A), R_g (B) and hSAS (C) of IAP (22–28) protofibrillar assembly in the absence (black) and in the presence of different intercalators (red: EtBr, blue: Dxr, green: Mtx and Chl: magenta). Red and blue arrows in (A) indicate the events of protofibril assembly destabilization by EtBr and Dxr respectively. (D–H) represents corresponding free energy profiles (kcal mol^{-1}) of IAP (22–28) protofibrillar assembly in the absence (D) and presence (E–H) of EtBr, Dxr, Mtx and Chl respectively. Corresponding lowest free energy conformations are shown as insets (grey cartoons) while intercalators are colored orange.



represented assemblies with higher gyration radius (R_g 1.17 nm) with phenanthridinium rings intercalated at site 2. Dxr induced mild deviations in packing of assembly (R_g 1.08 nm), attributed to engagement of single Dxr molecule at site 2 through its methyloxan unit while other Dxr molecules were bound at site 1. Notably, no such drift in acquired conformational space was observed for assembly in the presence Mtx and Chl, where binding was observed at site 1 alone (Fig. 4G and H). The lowest energy conformations corresponded to control-like compact assembly ($R_g \sim 1.01$ nm), although in these cases minimum disorder was induced in fibrils.

Molecular docking of intercalators to Lys and PrP

In Lys, preferred binding cavity for benzophenanthridine alkaloids and Chl had already been reported previously.⁴⁶ Interestingly, cluster analysis of binding poses post docking of other intercalators also displayed same preferred site for binding (ESI Fig. 8A†). For PrP, binding of intercalators were not restricted to a single fold as seen for Lys. We observed highest binding poses at different sites ($\alpha 2$ and $\alpha 3$ helices) of PrP (ESI Fig. 8B†). The highest docking free energies (kcal mol^{-1}) for binding of intercalators to Lys and PrP are summarized in ESI Table 2.†

Biophysical characterization of modulatory effects of intercalators

Motivated by the simulation estimates, we further extended our investigations to validate efficacy of intercalators against aggregation of IAP (22–28) to structured intermediates and modulation of its protofibril assembly through real-time *in vitro* aggregation assays. We were equally interested in testing if our computational estimates are generic in nature. This is important, since conformational transition of protein systems to their amyloid states rely on traversing through various kinetic and thermodynamic barriers, unlike in peptide systems where direct contributions from local interactions govern their spontaneous aggregation. The simulation predictions on interference mechanisms were also extended to experimental investigations against full length proteins, Lys and PrP. In all cases, the kinetics of amyloid formation was monitored using standard thioflavin T (ThT) fluorescence and static light scattering experiments (SLS).

Intercalators inhibit transitions of monomers to structured assemblies

ThT fluorescence exhibits a hyperchromic shift in emission maximum upon binding to β -rich fibrillar structures during amyloidogenesis.^{37,47,48} Traversing from lag and exponential to stationary phase, amyloidogenesis generally follows sigmoidal kinetics.^{49,50} In our case, reactions of proteins/peptides were prepared in the presence (increasing molar equivalents) and absence of compounds and were subjected to amyloid aggregation.

Under given aggregating conditions, all the four peptides/proteins showed gradual increase in ThT fluorescence over reaction periods (Fig. 5). In controls sets, the ThT fluorescence showed distinct lag phases of different duration (ESI Table 3†), with shortest observed for PrP.

Interestingly, in the presence of intercalators (regardless of the type), ThT yields remained persistently low at all ratios (2 : 1, 1 : 1 and 1 : 3) throughout the reaction time (Fig. 5E–P). ThT intensities failed to escalate exponentially (low I_{max} , maximum ThT intensity) suggesting total inhibition of nucleation dependent polymerization in a dose dependent manner (ESI Table 3†).

Interestingly, scattering for peptide-intercalator systems at lowest molar ratio (2 : 1) at the end of reaction period was moderately lower (~ 20 –30%) than corresponding controls, contrary to corresponding ThT yields. This indicated that compounds were effective in preventing formation of fibrillar assemblies (low ThT) by steering the aggregation of monomers towards large unstructured (high scattering) aggregates (ESI Fig. 9†). Accordingly at higher ratios (1 : 1 and 1 : 3), the systems displayed even lower scattering. The inhibitory control over the aggregation of peptide/protein monomers to fibrillar assemblies was established through TEM images of same systems where no fibrillar assemblies were observed (Fig. 6).

Intercalators arrest protofibrillar progression into mature fibrils

To validate our *in silico* predictions of inhibitory control of intercalators over the progression of aggregation pathway from pre-formed protofibrillar oligomers, the compounds were co-incubated independently at the mid-point of the exponential phases of individual systems (after 22 hours for IAP (22–28), 25 hours for AGel (186–192), 70 hours for Lys and 10 hours for PrP). Immediately after addition, the ThT intensity declined drastically to basal levels (Fig. 7). However, when the corresponding scattering profiles were analyzed, only marginal drop in intensities were observed (ESI Fig. 10†). After incubation with intercalators at this point, scattering of each system (at 2 : 1) failed to rise proportionally compared to controls and was nearly stagnant till the end of reaction period. This clearly suggested that intercalators obviated further assembly of protofibrillar oligomers to mature fibrils, thus arresting the pathway in a protofibrillar state. TEM images of these systems at the end of aggregation further confirmed the presence of protofibrillar species and absence of mature fibrils (ESI Fig. 11†).

Binding of intercalators marginally affects secondary and tertiary structure stability

Proteins may undergo structural changes upon exposure to exogenous factors as temperature, solvent environment which could result in alternate conformational states.^{10,51} For Lys and PrP, the observed anti-amyloidogenic effects could be attributed to induced local secondary or tertiary structure perturbations by intercalators. Thus, we tried to assess the effect of compounds on their native secondary and tertiary structures. In both proteins, steady state fluorescence titration with compounds (ESI Fig. 12A–H†) showed no apparent shifts around λ_{340} indicative of unaltered tertiary architecture. In Lys, secondary structural transitions were also absent, as indicated by overlapping CD spectra at all ratios (ESI Fig. 12I–L†). However, association of intercalators slightly induced secondary structure



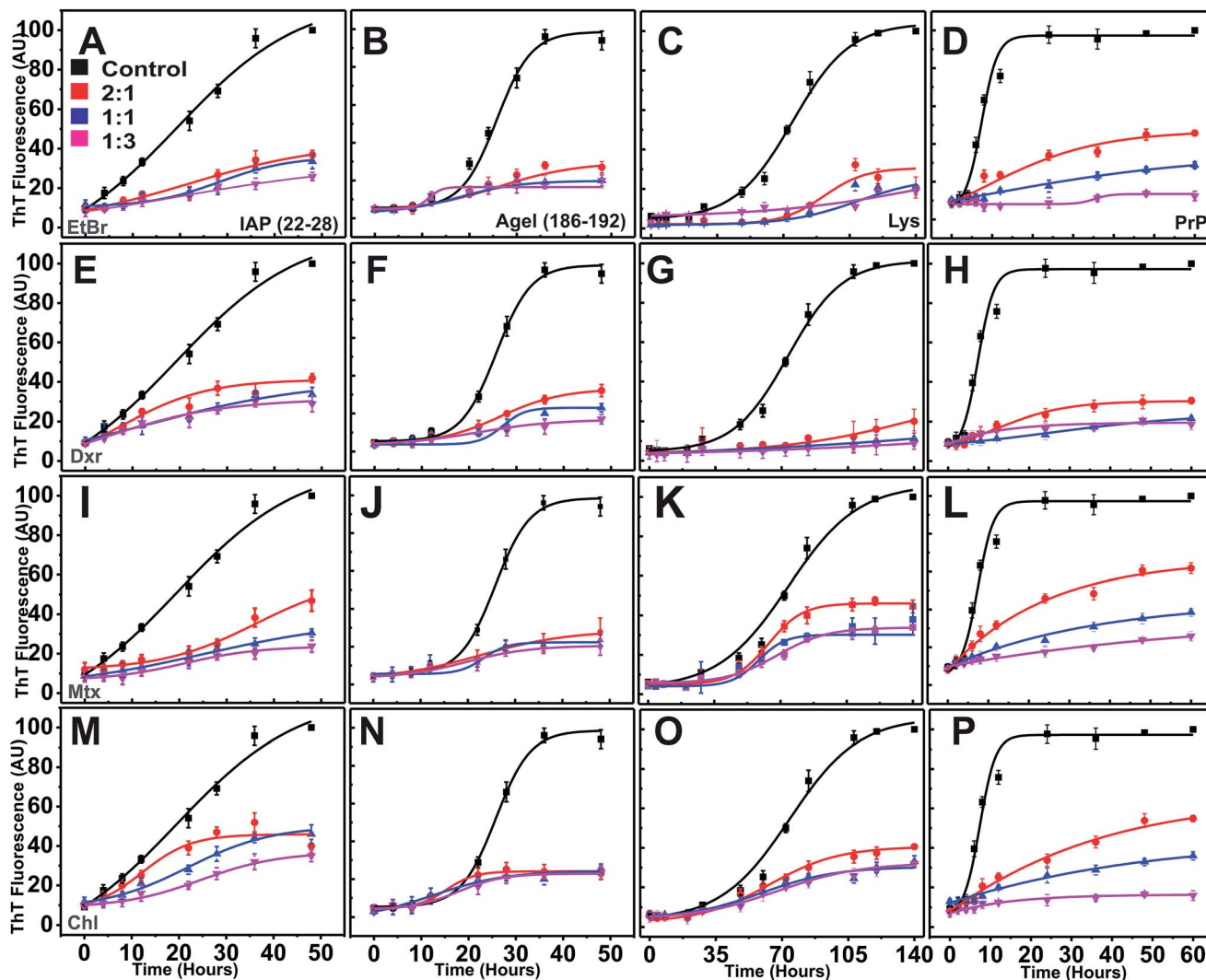


Fig. 5 Kinetics of protein/peptide amyloid formation. Normalized ThT fluorescence for aggregation kinetics of IAP (22–28) (100 μ M), AGel (186–192) (100 μ M), Lys (140 μ M) and PrP (30 μ M) (A–D) in the absence (black curves represents controls) and presence of EtBr at increasing P : D molar ratios (red 2 : 1, blue 1 : 1 and magenta 1 : 3) when the compounds were pre-incubated at initiation of lag phases (0 hour). Similarly, (E–H), (I–L) and (M–P) represents fibrillation kinetics in the presence of Dxr, Mtx and Chl respectively. Error bars in each data point indicate standard deviation.

perturbations in PrP (ESI Fig. 12M–8P[†]). Thermal melting studies of control and test systems (2 : 1 [P]/[D]) by monitoring λ_{340} and θ_{222} as function of temperature resulted in overlapping denaturation profiles (ESI Fig. 13A–D[†]). However, T_m calculated through first derivative in the presence of intercalators showed slight shift towards higher temperatures (2–3 $^{\circ}$ C) for tertiary structure transitions in Lys and PrP. For secondary structure transitions in PrP, T_m shifted slightly towards lower temperature (5–6 $^{\circ}$ C) compared to control in accord with induced secondary structure perturbations while it remained unaltered for Lys.

Intercalators specifically bind to monomeric/native peptides/proteins

Furthermore, it became essential to ascertain if different intercalators could actually bind to peptides/proteins. Association of intercalators to the aggregation prone regions (fully accessible in peptides while partially in proteins) could directly or indirectly

modulate their aggregation pathway by possibly blocking intermolecular interaction sites.⁴⁶ Fluorimetric titrations of intercalators with peptide/proteins provided clues on changes in microspace of intercalators upon association with monomers. Titration curves for each system are shown in ESI Fig. 14.[†] The dissociation constants for each system are summarized in ESI Table 4.[†] Micromolar range dissociation constants indicated strong association of intercalators to peptide/protein monomers resulting in strong PD complexes, putatively incapable of organizing into higher order structural aggregates.

Association of intercalators with pre-formed fibrils

The intercalators could arrest protofibrillar progression when added during exponential growth phase of each system indicating strong association between intercalators and protofibrils. Similar experiments were carried to verify the association between intercalators and fibrils of peptides/proteins. Both



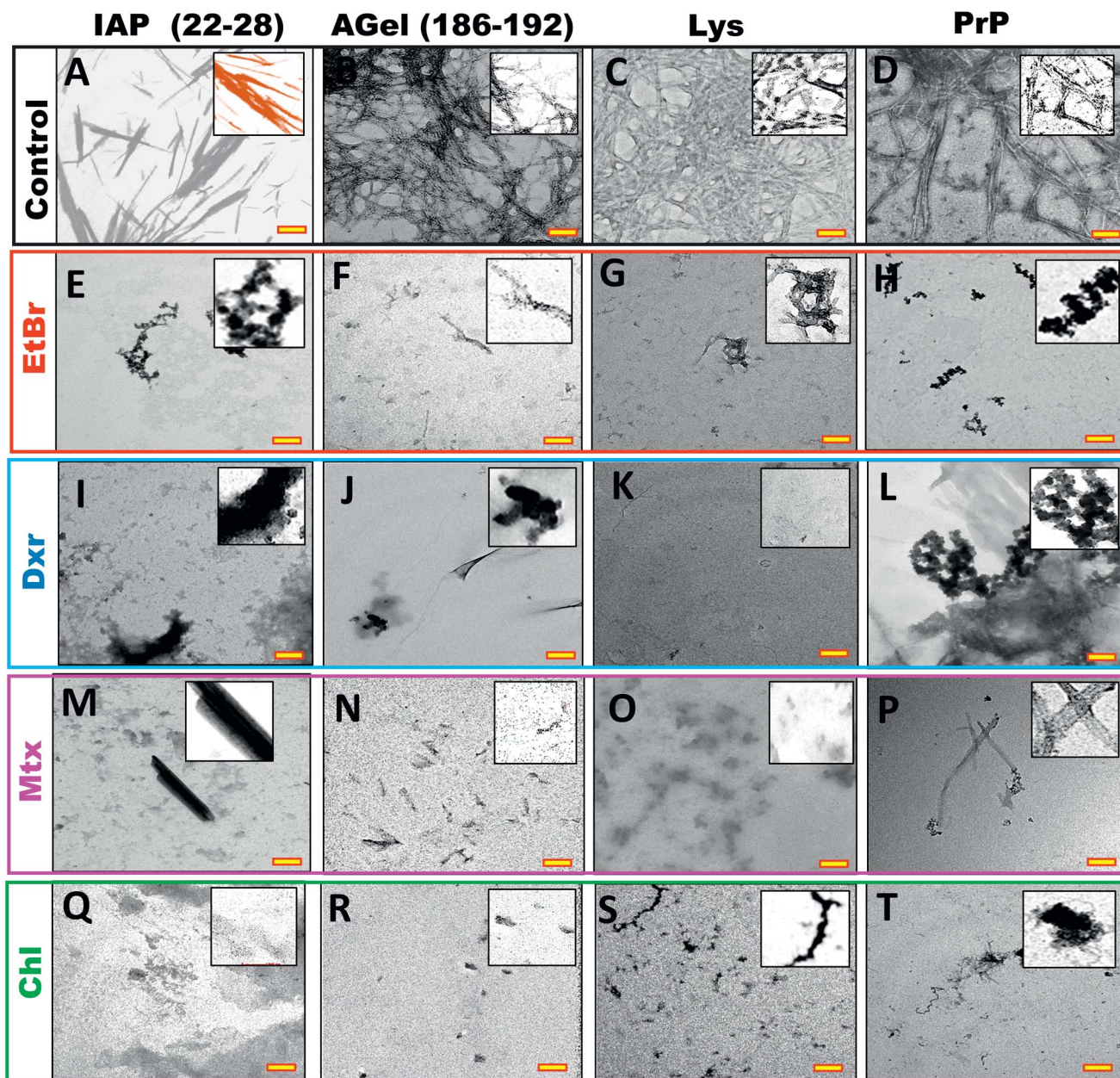


Fig. 6 TEM images of same systems in Fig. 7 (at 2 : 1 ratio), processed after 4 hours of their completion. End stage aggregates from fibrillation of IAP (22–28), AGel (186–192), Lys and PrP alone (A–D) and assembly process in the presence of EtBr, Dxr, Mtx and Chl (E–T). All intercalators showed complete inhibition of fibrillation independent of peptide/protein system. Scale bars (yellow insets) are 500 nm. Insets show magnified version of the sample.

hypo- and hyperchromic shifts in the fluorescence yield were visible for different compounds when titrated with corresponding pre-formed fibrils. Each trend represents distinct changes in the micro space of intercalators upon interacting with fibril assembly. Titration curves for each system were plotted as above (ESI Fig. 15[†]) and the dissociation constants are presented in ESI Table 5.[†]

Discussions

The application of many planar molecules has been extended as amyloid inhibitors where some of these molecules are proven to

reduce associated toxicities.^{28,29} In our study, we have attempted to gain insights into interference mechanisms of intercalators with the aggregation pathways of peptides and proteins, specifically emphasizing and establishing the role of hetero-aromatic associations.

Spontaneous aggregation of peptide monomers from IAP and gelsolin proteins to form structurally heterogeneous assemblies was established using μ s explicit solvent simulations. The simulation results agreed to initial dependency on solvation free energy of IAP (22–28) and AGel (186–192) monomers owing to high hydrophobicity (of 75% and 62% residues respectively)⁴⁵ to form initial collapsed state. Further optimal



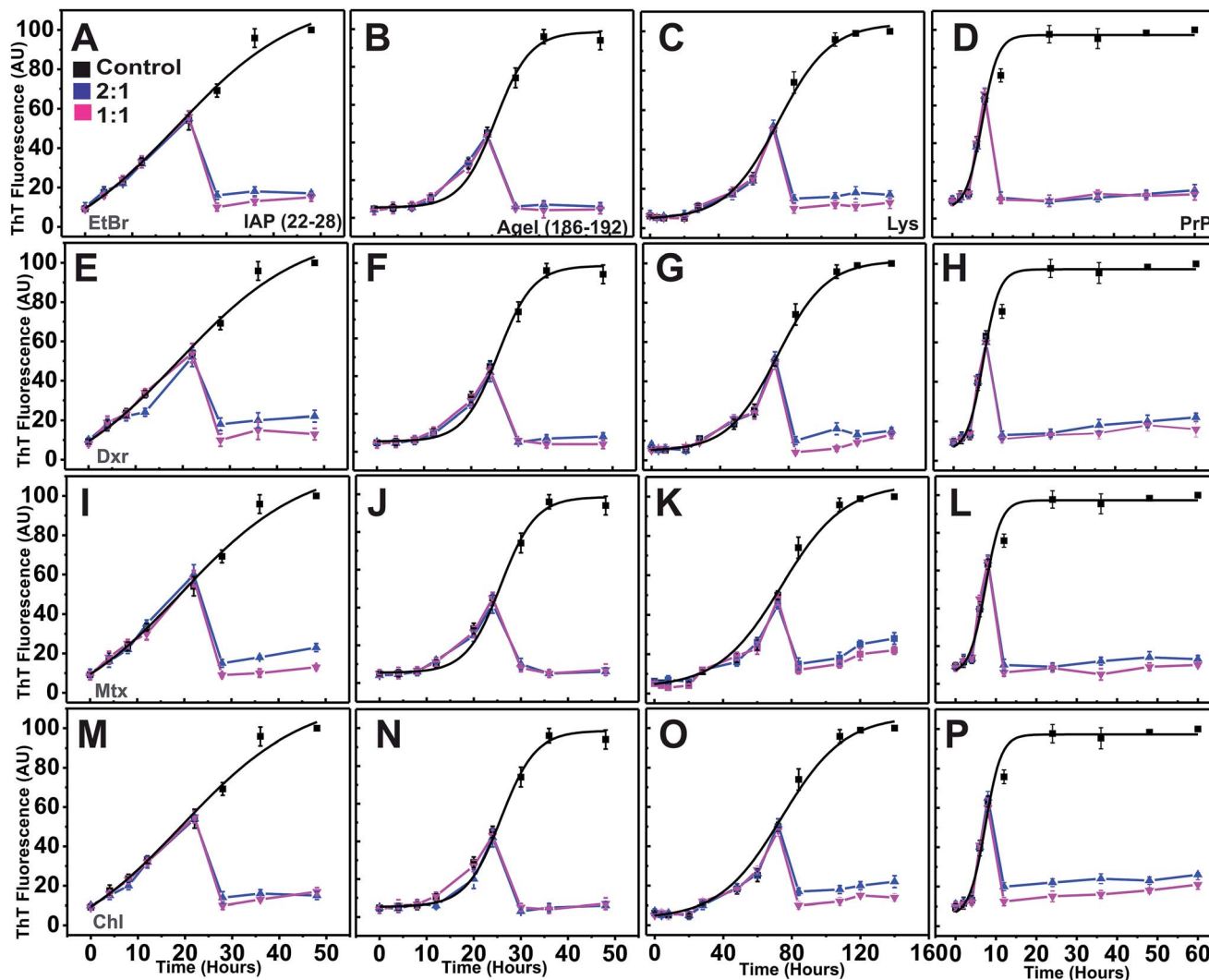


Fig. 7 Modulation of protofibril progression. Normalized ThT fluorescence for aggregation kinetics of IAP (22–28) (100 μ M), A β (186–192) (100 μ M), Lys (140 μ M) and PrP (30 μ M), (A–D) in the absence (black curves represents controls) and presence of EtBr at increasing P : D molar ratios (blue 2 : 1 and magenta 1 : 1) when the compounds were incubated during exponential growth phase of each system. Similarly, (E–H), (I–L) and (M–P) represents ThT kinetics in the presence of Dxr, Mtx and Chl respectively. Error bars in each data point indicate standard deviation.

structural reorganizations to partially structured species during the downhill process were reflected through incessant decline in dRMS with high increment in β -content during later stages of simulation. These observations complied with earlier experimental and theoretical reports documented on aggregation of short peptides.^{18,20,43,44,52} FEL projections indicated spontaneous association of monomers to form higher order ensembles whereas an early single large basin was observed in FEL of IAP (22–28) and A β (186–192) monomeric transitions into assembly in the presence of intercalators. Apparently, the kinetics of assembly formation in the presence of compounds was driven further owing to some topological advantage imparted by their planar groups. However, extraction of corresponding lowest energy conformation showed improper packing of monomers, terminating in non-amyloid oligomeric assemblies. Consequently, loss of inter-chain contacts was compensated by subsequent concurrent contacts between

monomers and compounds, particularly the stable hetero-aromatic associations with Phe residues. Binding poses of stable complexes obtained at the end of simulations clearly displayed hetero-aromatic associations between Phe's and planar groups of the compounds (ESI Fig. 7A–D†). Since the compounds spend most of the simulation period in associations with some Phe residues and their self-association, such hetero-aromatic associations could be accounted for modulation of aggregation pathways.

In vitro substantiation of simulation outcomes were highly corroborating and validated incompetency of IAP (22–28) and A β (186–192) monomers to form β -rich aggregates (ThT insensitive species) in the presence of intercalators. Similarly, emergence of stable disordered aggregates at the end of simulations was highly supported by SLS experiments where the corresponding scattering at 2 : 1 [peptide]/[D] was marginally lower than the respective controls. Binding affinity studies



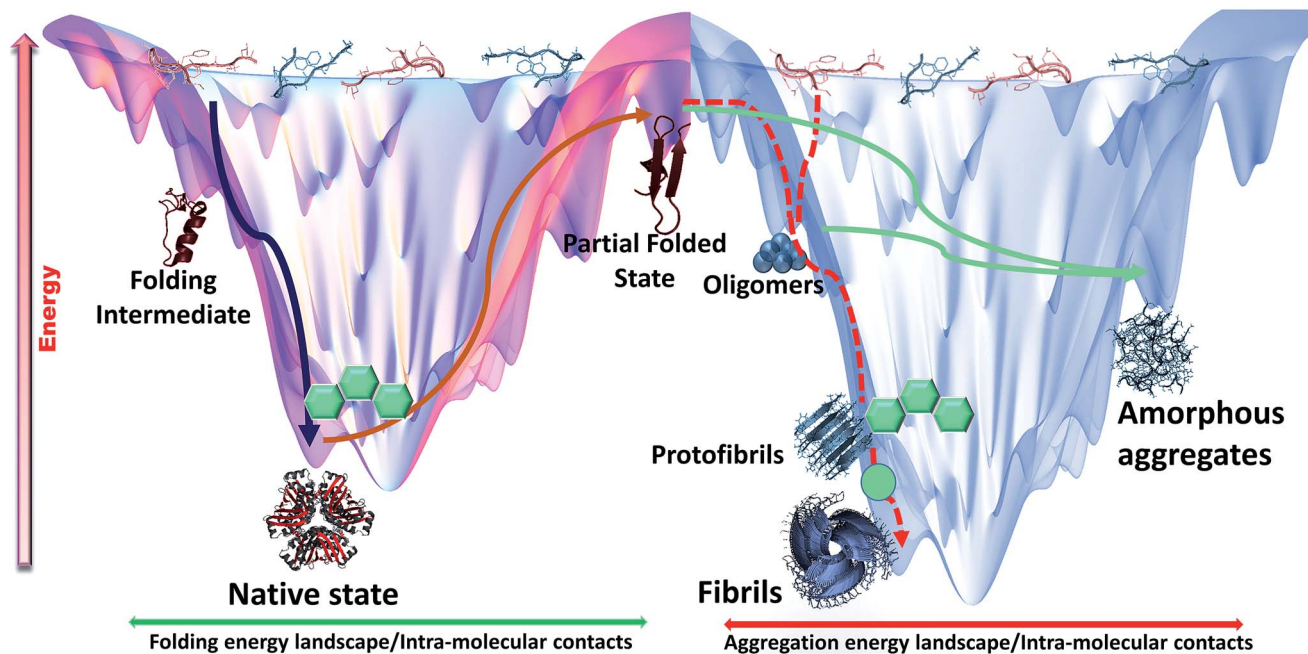


Fig. 8 The competing folding and aggregation free energy landscapes. The colored free energy funnel corresponds to downhill folding (blue arrow) of polypeptides to native state via transient folding intermediates, dominated by intra-molecular interactions. On the right, the pale blue energy funnel corresponds aggregation landscape dominated by inter-molecular interactions. Destabilization of native states or non-native folding (orange arrow) may lead to their entrapment as high energy partially folded states. These states further funnel down (red arrows) to either amorphous (disordered) or amyloid (ordered) states. The fibrillar arrangement of amyloids caters to lowest energy conformation formed through associations of basic structural unit, protofibrils. The intercalators (colored green) upon interactions with native states drive the aggregation pathway towards unstructured amorphous states (light green dashed arrows), thus impeding formation of toxic intermediates. Alternatively, these also interact with pre-formed protofibrils interfering with their assembly to mature fibrils. The energy landscapes are constructed using modified code for generating funnels from Oas Lab (Departments of Biochemistry & Chemistry Duke University, USA).

showed that the peptide monomers were initially capable of forming stable (PD) complexes with intercalators. These inhibitory effects were reciprocated by the Lys and PrP as well. We reasoned that having significant binding affinities (μM range), the intercalators may confer additional stabilizing effect to the protein's native structure, resisting their amyloidogenic conversion. However, thermal denaturation studies ruled out this possibility and led us to propose that along the reaction pathway, these intercalators bind to core amyloidogenic regions (directly in case of peptide amyloidogenesis and indirectly after partial unfolding of proteins) and steering the pathway towards amyloid incompetent states.

The observed interference effects of intercalators could be explained by two mechanisms; hetero-association of intercalators with peptide monomers and self-association of intercalators. The hetero-association between intercalators and peptide monomers can occlude the amyloid-like interactions by blocking the potential inter-chain free energy surfaces, crucial for assembly process.⁵³ This was indeed supported through observed low dissociation constants of peptides with intercalators. Furthermore in our simulations, we observed self-association of intercalators with concurrent recruitment of peptide monomers to the opposite end of their assembly, possibly providing stable scaffolds for hetero-molecular interactions. These associations were coherent with previously reported evidences on macromolecule dependent/independent

association of ligands.^{54–59} This super assembly can be represented as monomer-(intercalator)_n-monomer, where n represents number of compound monomers involved in self-assembly. Thus, it could be conceived from the observed facts that compounds were capable of intercalating in between monomeric peptides during the association events itself, driving the aggregation pathway towards alternate assemblies, incompetent to form 'on-pathway' pre-fibrillar oligomer assembly. On the other hand, in case of Lys, binding preferences of EtBr, Dxr and Mtx also incorporates amyloidogenic stretches which could be directly or indirectly implicated in averting its amyloidogenesis (ESI Fig. 8†). Interestingly, the preferential modes for PrP suggested the possibility of hetero-aromatic interactions with crucial aromatic residues at the $\alpha 2$ - $\alpha 3$ helices. The unfolding of these helices have been shown to initiate PrP fibrillogenesis and the intercalators mostly stabilize this crucial tertiary scaffold⁶⁰ (ESI Fig. 13B†).

The intercalators actively induced remodeling of the pre formed protofibril assembly. From the structural perspective, protofibrils represent basic building units for elongation to mature amyloid fibrils. Coarse grained simulations on IAP (20–29) fibril formation and its elongation, as reported by Sorensen *J. et al.*, depicts these associations mediated through solvent exposed residues at all interfaces, promoting inter molecular contacts with other assemblies in system.⁵³ On the other hand, binding of Congo red and phenol red are earlier shown to



induce subtle structural changes to the protofibrils formed by the core amyloid stretches of yeast prion protein Sup35 and hIAPP.^{61,62} However, owing to smaller simulation time scales (20 ns) in these studies, no gross destabilization of protofibrils was observed.

However, in our studies EtBr and Dxr associated with the IAP (22–28) protofibrils both longitudinally and laterally (S1 and S2 sites respectively). These associations induced structural perturbations at later stages of simulations as depicted through higher presentation of hydrophobic side chains to the solvent (Fig. 4C). However, Mtx and Chl interactions were only confined to exposed residues at fibril surface which further resulted in even lower solvent exposure of hydrophobic residues than control, thus potentially blocking the longitudinal interaction sites. Both types of binding could interfere with further assembly of individual protofibrils, resulting in null/stunted fibril growth. Speculations for interference with protofibril elongation in the presence of intercalators were substantially supported by scattering and ThT profiles. The observed stagnancy in scattering after incubation of compounds at time intervals corresponding to dominant protofibril population in all systems (ESI Fig. 10†) strongly indicated towards ‘freezing’ the elongation of protofibrils to mature fibrils. Further, a sharp drop in corresponding ThT yields validated S1 as preferred site for intercalator–fibril interactions that plausibly prevents fibril growth. Recent work on Lys and insulin amyloids⁶³ showed fibril surface as single preferred site for ThT binding, accounting for its enhanced fluorescence upon association. Drawing parallels, ThT molecules would have been incapable of displacing the intercalators from their original binding site resulting in low quantum yields. This was indeed supported by apparently high binding affinities of intercalators with fibril assemblies (ESI Table 4†). The corresponding scattering intensities remained nearly constant till the end of reactions indicating stalling of amyloid progression from protofibrils by the intercalators. The possibility of an artefact arising due to marginal change in scattering in peptide systems is negated by the observance of substantial change in scattering profiles for protein samples.

The simulation projections provided important mechanistic insights about interaction and interference of intercalators with amyloidogenic pathways (Fig. 8). Simulation results were conclusively proven through *in vitro* experiments which suggested dominant inhibitory role of DNA intercalators during amyloid assembly process, facilitated predominantly through aromatic associations with native monomers or along their aggregation process. However, the current approach to gain mechanistic insights could be further supported by parallel replica exchange molecular dynamics (REMD) studies to allow more coverage of conformational space and enhanced sampling efficiency. Recent reports on REMD studies on cross interactions of p53 core fragment and its aggregation rescue mutant⁶⁴ and of A β _{1–42} and hIAPP_{1–37} polypeptides⁶⁵ showed formation of diverse conformational assemblies within given time scale. Such detail might have been overlooked if conventional MD approach is exclusively used. Similarly, it is possible that other potential intermediates and terminal assemblies in the

presence of intercalators were missed out when an all-atomistic simulation is used. These unaccounted intermediates further needs to be explored through more extensive computational and experimental validations. Through multi-dimensional approaches, this study attempts to understand complex hetero-molecular landscape underlying anti-amyloid potential of planar molecules studied so far.

Materials and methodologies

Molecular dynamics system and simulations

Simulation setup. All simulations were carried using GRO-MACS 5.1 software package under gromos 54A7 force field.^{66–68} The initial structures of test compounds were imported from PubChem Compound Database while the topology and other parameters were assigned using ATB topology builder.^{69,70} To study the interference of intercalators with amyloidogenic pathways of peptide and proteins, two different types of systems were setup; aggregation of peptide monomers to form protofibril like oligomers and effects of intercalators on pre-formed protofibrillar assembly. In first system, eight pre-generated random conformation monomers of IAP (22–28) and AGel (186–192) were simulated separately, corresponding to their experimental aggregation conditions described elsewhere,^{20,37} without applying backbone dihedral or distance constraints. In identical setups, simulations were carried out in the presence of intercalators to assess their ability in modulating assembly formation by respective peptides. Briefly, the monomers of IAP (22–28) and AGel (186–192) were placed in random orientation in individual cubical boxes (ESI Fig. 1A†) to achieve a final concentration of ~66 mM. Subsequently, in separate setups, eight monomers of EtBr, Dxr, Mtx and Chl were placed at random positions and orientations in each cubical box such that the peptide (P) to compound (D) molar ratio remains <1. Lower molar ratio was preferred to avoid possible bias owing to overcrowding of intercalators. In each setup, organization of monomers to a supra-molecular assembly, analogous to reference structure was mapped. The reference frame represented an octameric protofibril assembly formed from their respective monomers. For IAP (22–28) monomer simulations (1 μ s for each system), pre-available protofibrillar structure (PDB id 2kib) served as reference frame. However for AGel (186–192) simulations (500 ns each), the octameric protofibril was initially constructed based on 2kib atomic coordinates and subsequently simulated for 500 ns prior using as reference frame.

To obtain binding free energies of compounds with the 2kib assembly, the compounds were initially docked with fibrillar octameric structure at 1:1 ratio (octamer assembly : compound) by using inbuilt Site Finder application in Molecular Operating Environment v2010.12 (MOE).⁷¹ Protocols for site identification and free energy calculations for docking to fibril assembly and proteins are detailed elsewhere.⁷²

For protofibril simulations (1 μ s each), studies were restricted to hIAPP (PDB id 2KIB) where its octameric assembly alone was positioned at the center of a box with same dimensions as used earlier. Subsequently, in each separate setup, five molecules of test compounds were positioned randomly in the



box. This alternate system was employed to reduce the bias originating from pre-defined conformations of ligand, docked to octamer assembly. Overall, a cumulative of 12.5 μ s all atom MD simulations were carried out. Each system was explicitly solvated with appropriate amount of SPC water molecules and ions added to maintain electroneutrality at 0.1 M NaCl concentration equivalent to *in vitro* system setup for carrying IAP (22–28) and AGel (186–192) amyloidogenesis. Subsequently, post energy minimization of system by steepest descent scheme, 1 ns equilibration using an isothermal–isobaric and isochoric–isothermal ensemble was performed respectively. PME was used to treat long range electrostatics with short range interactions at cutoff radius of 10 Å for both coulomb and van der Waal potentials. System temperature (310 K) and pressure (1 bar) was kept constant using velocity rescale and Berendsen barostat respectively. Finally, the velocities were generated by solving Newton's equations of motion with 2 fs time step using Leap frog integrator. The coordinates were recorded at each 5 ps interval. The resulting trajectories were analyzed by inbuilt gromacs tools while the images were created using PyMol package.⁷³

Chemicals

The IAP (22–28) and AGel (186–192) peptides were custom synthesized and purchased from BioCell Corporation (India) as 99% pure lyophilized powder. Chicken egg-white lysozyme and all chemicals were purchased from Sigma Chemical Co. (St. Louis, MO). Human PrP was purified as per previous reports.⁷⁴

Preparation and characterization of amyloid fibrils

IAP (22–28) and AGel (186–192) amyloidogenesis. Stock solutions of each peptide was made in 100% DMSO. For IAP (22–28), amyloidogenesis was initiated by incubating 100 μ M peptide at 37 °C in 50 mM HEPES, pH 7.6 with constant shaking at 300 rpm.³⁶ AGel (186–192) amyloidogenesis was carried in sodium acetate, pH 4.0 with 100 mM NaCl at 37 °C without shaking.³⁷

Lys and PrP amyloidogenesis. For Lys amyloidogenesis, the reaction mixture contained 140 μ M Lys at 37 °C in acetate buffer, pH 7.2 with continuous agitation at 500 rpm. For PrP the reaction set up contained 30 μ M PrP in sodium phosphate buffer, pH 7.4 with continuous agitation at 330 rpm.

Amyloidogenesis in the presence of intercalators/compounds. Aggregation reactions were set up at protein/peptide [P] : compound [D] ratios of 2 : 1, 1 : 1 and 1 : 3 and monitored for ThT fluorescence. For assessing the effect of compounds on pre-formed protofibrils, compounds were added to the protofibrillar stage (mid-exponential phase of the ThT curve) at [P] : [D] ratios of 2 : 1 and 1 : 1. All measurements were carried out in triplicates wherein error bars represent standard deviation of each data point.

Fluorescence and CD measurements. Fluorimetric assays and determinations were performed using Perkin-Elmer LS55 spectrofluorimeter at 37 °C.

Thioflavin T (ThT) binding assays. At each sampling, 90 μ L of 10 μ M ThT in phosphate buffered saline (PBS), pH 7.4 was

mixed with 10 μ L of samples and were incubated for 15 minutes prior to fluorescence measurement. Negative controls of ThT incubated with compounds were setup in similar manner for background corrections. The emission/excitation wavelengths were set at 485/450 nm and slits at 10/5 respectively. The data was fitted according to earlier reports using following equation:

$$y = y_0 - y_{\max} / \left(1 + e^{-k \left(t - t_{\frac{1}{2}} \right)} \right)$$

where, y_0 and y indicates ThT fluorescence at $t = 0$ and any time (t) and y_{\max} is maximum ThT fluorescence. Lag phase was calculated using $y_{\frac{1}{2}} - 2/k$.

Scattering assays. Static light scattering (SLS) of the samples were recorded with both the excitation and emission wavelengths kept at 350 nm with slits at 5/5.

Intrinsic fluorescence assays. Samples at P : D ratios 2 : 1, 1 : 1 and 1 : 3 were incubated for 4 hours prior to fluorescence scans between 300 nm to 400 nm with excitation at 290 nm. Excitation and emission slits were set at 5/5. Thermal denaturation assays at P : D (2 : 1) were performed by gradual heating from 20 °C to 90 °C with ramping rate of 1° per minute.

Far UV CD. For far UV CD spectroscopy samples were taken in a 2 mm path length quartz cuvette and scanned from 250–200 nm at scan rate of 200 nm per minute in a Jasco CD spectropolarimeter (Japan). Thermal denaturation assays were similarly performed to assess effect of intercalators on secondary structure transitions. Change in ' θ_{222} ' was monitored against temperature with ramping rate at 1° per minute.

Intrinsic drug fluorescence assays. Changes in the micro-environment of the compounds upon binding to monomers or to protofibrils were probed by fluorometric titrations. Each compound taken at 2 μ M concentration in a 1 cm path length quartz cuvette was titrated against increasing concentrations of protein or peptide monomers. Titrations with increasing concentrations of pre-formed fibrils of each protein/peptide were carried out in a similar manner. Briefly, proteins/peptides subjected to amyloidogenic conditions in respective buffers (as mentioned earlier) were harvested by centrifugation at 14 000 rpm for 1 hour after incubation for 60 h. The pellet was suspended in 50 μ L respective buffer and sonicated for 5 minutes. The sonicated samples were resuspended in 450 μ L of respective buffer before use in the titration experiments. All scans were recorded with excitation/emission slits set at 5/10 nm. The excitation wavelength for EtBr and Dxr was set at 485 nm while for Mtx and Chl it was kept at 660 nm and 420 nm, respectively. Change in fluorescence intensity was plotted against increasing concentration of protein/peptide. Dissociation constant (K_d) was obtained by using GraphPad Prism 5 (GraphPad Software Inc., San Diego, CA) employing nonlinear regression least square curve fitting analysis using single site specific model as:

$$Y = B_{\max} \times X / (K_d + X)$$

where, B_{\max} represents maximum specific binding and K_d is the equilibrium dissociation constant.



Morphology of aggregates. TEM (Transmission Electron Microscopy) was used to visualize aggregation of peptides and proteins in the absence and presence of intercalators. Samples for analysis were withdrawn post 4 hours from the completion of saturation phase of each system. Briefly, each specimen was prepared by depositing 5 μL sample over carbon coated copper grids and incubated for 5 min. The grids were then rinsed twice with 10 μL Milli-Q water followed by their staining with 2% uranyl acetate for 2 min. Air dried grids were examined using FEI Tecnai TF20 operated at maximum accelerating voltage of 200 kV.

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References

- W. M. Tay, D. Huang, T. L. Rosenberry and A. K. Paravastu, *J. Mol. Biol.*, 2013, **425**, 2494–2508.
- E. C. Mader Jr, R. El-Abassi, N. R. Villemarette-Pittman, L. Santana-Gould, P. W. Olejniczak and J. D. England, *Neurol. Int.*, 2013, **5**, e1.
- P. Trepte, N. Stempel and E. E. Wanker, *Essays Biochem.*, 2014, **56**, 167–180.
- W. J. Du, J. J. Guo, M. T. Gao, S. Q. Hu, X. Y. Dong, Y. F. Han, F. F. Liu, S. Jiang and Y. Sun, *Sci. Rep.*, 2015, **5**, 7992.
- F. Chiti and C. M. Dobson, *Annu. Rev. Biochem.*, 2006, **75**, 333–366.
- M. Sunde, L. C. Serpell, M. Bartlam, P. E. Fraser, M. B. Pepys and C. C. Blake, *J. Mol. Biol.*, 1997, **273**, 729–739.
- S. Kumar and J. B. Udgaonkar, *J. Mol. Biol.*, 2009, **385**, 1266–1276.
- C. G. Glabe and R. Kaye, *Neurology*, 2006, **66**, S74–S78.
- C. Goldsbury, J. Kistler, U. Aebi, T. Arvinte and G. J. Cooper, *J. Mol. Biol.*, 1999, **285**, 33–39.
- J. C. Rochet and P. T. Lansbury Jr, *Curr. Opin. Struct. Biol.*, 2000, **10**, 60–68.
- R. Carrota, M. Manno, D. Bulone, V. Martorana and P. L. San Biagio, *J. Biol. Chem.*, 2005, **280**, 30001–30008.
- G. T. Dolphin, P. Dumy and J. Garcia, *Angew. Chem.*, 2006, **45**, 2699–2702.
- M. Cheon, I. Chang, S. Mohanty, L. M. Luheshi, C. M. Dobson, M. Vendruscolo and G. Favrin, *PLoS Comput. Biol.*, 2007, **3**, 1727–1738.
- S. Auer, F. Meersman, C. M. Dobson and M. Vendruscolo, *PLoS Comput. Biol.*, 2008, **4**, e1000222.
- A. De Simone, A. Dhulesia, G. Soldi, M. Vendruscolo, S. T. Hsu, F. Chiti and C. M. Dobson, *Proc. Natl. Acad. Sci. U. S. A.*, 2011, **108**, 21057–21062.
- A. R. Dinner, A. Sali, L. J. Smith, C. M. Dobson and M. Karplus, *Trends Biochem. Sci.*, 2000, **25**, 331–339.
- A. Srivastava, P. Arya, S. Goel, B. Kundu, P. Mishra and A. Fnu, *PLoS One*, 2015, **10**, e0127011.
- C. Wu, H. Lei and Y. Duan, *Biophys. J.*, 2005, **88**, 2897–2906.
- D. Zanuy, B. Ma and R. Nussinov, *Biophys. J.*, 2003, **84**, 1884–1894.
- L. H. Tu and D. P. Raleigh, *Biochemistry*, 2013, **52**, 333–342.
- K. E. Marshall, K. L. Morris, D. Charlton, N. O'Reilly, L. Lewis, H. Walden and L. C. Serpell, *Biochemistry*, 2011, **50**, 2061–2071.
- M. L. Waters, *Curr. Opin. Chem. Biol.*, 2002, **6**, 736–741.
- P. Yakovchuk, E. Protozanova and M. D. Frank-Kamenetskii, *Nucleic Acids Res.*, 2006, **34**, 564–574.
- S. Stefansson, D. L. Adams and C. M. Tang, *BioTechniques*, 2012, **52**, 1–6.
- Y. Porat, A. Abramowitz and E. Gazit, *Chem. Biol. Drug Des.*, 2006, **67**, 27–37.
- S. Bastianetto, S. Krantic and R. Quirion, *Mini-Rev. Med. Chem.*, 2008, **8**, 429–435.
- A. M. Wardley, G. C. Jayson, D. J. Goldsmith, M. C. Venning, P. Ackrill and J. H. Scarffe, *Br. J. Cancer*, 1998, **78**, 774–776.
- G. Merlini, E. Ascari, N. Amboldi, V. Bellotti, E. Arbustini, V. Perfetti, M. Ferrari, I. Zorzoli, M. G. Marinone, P. Garini, et al., *Proc. Natl. Acad. Sci. U. S. A.*, 1995, **92**, 2959–2963.
- R. Colombo, A. Carotti, M. Catto, M. Racchi, C. Lanni, L. Verga, G. Caccialanza and E. De Lorenzi, *Electrophoresis*, 2009, **30**, 1418–1429.
- K. Augutis, M. Axelsson, E. Portelius, G. Brinkmalm, U. Andreasson, M. K. Gustavsson, C. Malmstrom, J. Lycke, K. Blennow, H. Zetterberg and N. Mattsson, *Mult. Scler.*, 2013, **19**, 543–552.
- A. Mukherjee, R. Lavery, B. Bagchi and J. T. Hynes, *J. Am. Chem. Soc.*, 2008, **130**, 9747–9755.
- A. Banerjee, J. Singh and D. Dasgupta, *J. Fluoresc.*, 2013, **23**, 745–752.
- S. Lahiri, T. Takao, P. G. Devi, S. Ghosh, A. Ghosh, A. Dasgupta and D. Dasgupta, *BioMetals*, 2012, **25**, 435–450.
- S. Lahiri, P. G. Devi, P. Majumder, S. Das and D. Dasgupta, *J. Phys. Chem. B*, 2008, **112**, 3251–3258.
- D. Reha, M. Kabelac, F. Ryjacek, J. Sponer, J. E. Sponer, M. Elstner, S. Suhai and P. Hobza, *J. Am. Chem. Soc.*, 2002, **124**, 3366–3376.
- A. Srivastava, S. Sharma, S. Sadanandan, S. Gupta, J. Singh, S. Gupta, V. Haridas and B. Kundu, *Biochem. J.*, 2016, DOI: 10.1042/BCJ20160737.
- P. Arya, A. Srivastava, S. V. Vasaikar, G. Mukherjee, P. Mishra and B. Kundu, *ACS Chem. Neurosci.*, 2014, **5**, 982–992.
- F. Bemporad, N. Taddei, M. Stefani and F. Chiti, *Protein Sci.*, 2006, **15**, 862–870.
- J. T. Nielsen, M. Bjerring, M. D. Jeppesen, R. O. Pedersen, J. M. Pedersen, K. L. Hein, T. Vosegaard, T. Skrydstrup, D. E. Otzen and N. C. Nielsen, *Angew. Chem.*, 2009, **48**, 2118–2121.
- J. P. Solomon, I. T. Yonemoto, A. N. Murray, J. L. Price, E. T. Powers, W. E. Balch and J. W. Kelly, *Biochemistry*, 2009, **48**, 11370–11380.
- L. A. Morozova-Roche, J. Zurdo, A. Spencer, W. Noppe, V. Receveur, D. B. Archer, M. Joniau and C. M. Dobson, *J. Struct. Biol.*, 2000, **130**, 339–351.



- 42 M. Jeffrey, G. McGovern, N. Makarava, L. Gonzalez, Y. S. Kim, R. G. Rohwer and I. V. Baskakov, *Neuropathol. Appl. Neurobiol.*, 2014, **40**, 296–310.
- 43 D. Matthes, V. Daebel, K. Meyenberg, D. Riedel, G. Heim, U. Diederichsen, A. Lange and B. L. de Groot, *J. Mol. Biol.*, 2014, **426**, 362–376.
- 44 D. Matthes, V. Gapsys, V. Daebel and B. L. de Groot, *PLoS One*, 2011, **6**, e19129.
- 45 J. Kyte and R. F. Doolittle, *J. Mol. Biol.*, 1982, **157**, 105–132.
- 46 C. Jash, P. Basu, P. V. Payghan, N. Ghoshal and G. S. Kumar, *Phys. Chem. Chem. Phys.*, 2015, **17**, 16630–16645.
- 47 M. Biancalana and S. Koide, *Biochim. Biophys. Acta*, 2010, **1804**, 1405–1412.
- 48 H. LeVine III, *Protein Sci.*, 1993, **2**, 404–410.
- 49 M. Bartolini, C. Bertucci, M. L. Bolognesi, A. Cavalli, C. Melchiorre and V. Andrisano, *ChemBioChem*, 2007, **8**, 2152–2161.
- 50 D. Hamada and C. M. Dobson, *Protein Sci.*, 2002, **11**, 2417–2426.
- 51 J. W. Kelly, *Curr. Opin. Struct. Biol.*, 1996, **6**, 11–17.
- 52 S. M. Tracz, A. Abedini, M. Driscoll and D. P. Raleigh, *Biochemistry*, 2004, **43**, 15901–15908.
- 53 J. Sorensen, X. Periolo, K. K. Skeby, S. J. Marrink and B. Schiott, *J. Phys. Chem. Lett.*, 2011, **2**, 2385–2390.
- 54 K. Karatasos, *J. Phys. Chem. B*, 2013, **117**, 2564–2575.
- 55 M. Menozzi, L. Valentini, E. Vannini and F. Arcamone, *J. Pharm. Sci.*, 1984, **73**, 766–770.
- 56 M. Enache and E. Volanschi, *Rev. Roum. Chim.*, 2010, **55**, 255–262.
- 57 D. B. Davies, D. A. Veselkov, M. P. Evstigneev and A. N. Veselkov, *J. Chem. Soc., Perkin Trans. 2*, 2001, 61–67, DOI: 10.1039/B007042o.
- 58 M. P. Evstigneev, V. P. Evstigneev and D. B. Davies, *J. Mol. Struct.*, 2006, **784**, 162–168.
- 59 B. P. Roques, J. Barbet and J. B. Lepecq, *Comptes Rendus Hebdomadaires Des Seances De L Academie Des Sciences Serie D*, 1976, **283**, 1453–1455.
- 60 M. Adrover, K. Pauwels, S. Prigent, C. de Chiara, Z. Xu, C. Chapuis, A. Pastore and H. Rezaei, *J. Biol. Chem.*, 2010, **285**, 21004–21012.
- 61 C. Wu, H. Lei, Z. Wang, W. Zhang and Y. Duan, *Biophys. J.*, 2006, **91**, 3664–3672.
- 62 C. Wu, Z. Wang, H. Lei, W. Zhang and Y. Duan, *J. Am. Chem. Soc.*, 2007, **129**, 1225–1232.
- 63 V. Ivancic, O. Ekanayake and N. Lazo, *ChemPhysChem*, 2016, DOI: 10.1002/cphc.201600246.
- 64 J. Lei, R. Qi, G. Wei, R. Nussinov and B. Ma, *Phys. Chem. Chem. Phys.*, 2016, **18**, 8098–8107.
- 65 M. Zhang, R. Hu, H. Chen, X. Gong, F. Zhou, L. Zhang and J. Zheng, *J. Chem. Inf. Model.*, 2015, **55**, 1628–1639.
- 66 Z. Lin and W. F. van Gunsteren, *J. Comput. Chem.*, 2013, **34**, 2796–2805.
- 67 W. Huang, Z. Lin and W. F. van Gunsteren, *J. Chem. Theory Comput.*, 2011, **7**, 1237–1243.
- 68 N. Schmid, A. P. Eichenberger, A. Choutko, S. Riniker, M. Winger, A. E. Mark and W. F. van Gunsteren, *Eur. Biophys. J.*, 2011, **40**, 843–856.
- 69 K. B. Koziara, M. Stroet, A. K. Malde and A. E. Mark, *J. Comput.-Aided Mol. Des.*, 2014, **28**, 221–233.
- 70 A. K. Malde, L. Zuo, M. Breeze, M. Stroet, D. Poger, P. C. Nair, C. Oostenbrink and A. E. Mark, *J. Chem. Theory Comput.*, 2011, **7**, 4026–4037.
- 71 S. Vilar, G. Cozza and S. Moro, *Curr. Top. Med. Chem.*, 2008, **8**, 1555–1572.
- 72 J. Singh, A. Srivastava, P. Jha, K. K. Sinha and B. Kundu, *Mol. Biosyst.*, 2015, **11**, 1887–1896.
- 73 Schrodinger, LLC, unpublished work.
- 74 B. Kundu, N. R. Maiti, E. M. Jones, K. A. Surewicz, D. L. Vanik and W. K. Surewicz, *Proc. Natl. Acad. Sci. U. S. A.*, 2003, **100**, 12069–12074.

