

Article

Alzheimer's Protective A2T Mutation Changes the Conformational Landscape of the A β_{1-42} Monomer Differently Than Does the A2V Mutation

Payel Das,^{1,*} Brian Murray,² and Georges Belfort²¹Soft Matter Theory and Simulations Group, Computational Biology Center, IBM Thomas J. Watson Research Center, Yorktown Heights, New York; and ²Howard P. Isermann Department of Chemical and Biological Engineering and Center for Biotechnology and Interdisciplinary Studies, Rensselaer Polytechnic Institute, Troy, New York

ABSTRACT The aggregation of amyloid- β (A β) peptides plays a crucial role in the etiology of Alzheimer's disease (AD). Recently, it has been reported that an A2T mutation in A β can protect against AD. Interestingly, a nonpolar A2V mutation also has been found to offer protection against AD in the heterozygous state, although it causes early-onset AD in homozygous carriers. Since the conformational landscape of the A β monomer is known to directly contribute to the early-stage aggregation mechanism, it is important to characterize the effects of the A2T and A2V mutations on A β_{1-42} monomer structure. Here, we have performed extensive atomistic replica-exchange molecular dynamics simulations of the solvated wild-type (WT), A2V, and A2T A β_{1-42} monomers. Our simulations reveal that although all three variants remain as collapsed coils in solution, there exist significant structural differences among them at shorter timescales. A2V exhibits an enhanced double-hairpin population in comparison to the WT, similar to those reported in toxic WT A β_{1-42} oligomers. Such double-hairpin formation is caused by hydrophobic clustering between the N-terminus and the central and C-terminal hydrophobic patches. In contrast, the A2T mutation causes the N-terminus to engage in unusual electrostatic interactions with distant residues, such as K16 and E22, resulting in a unique population comprising only the C-terminal hairpin. These findings imply that a single A2X (where X = V or T) mutation in the primarily disordered N-terminus of the A β_{1-42} monomer can dramatically alter the β -hairpin population and switch the equilibrium toward alternative structures. The atomistically detailed, comparative view of the structural landscapes of A2V and A2T variant monomers obtained in this study can enhance our understanding of the mechanistic differences in their early-stage aggregation.

INTRODUCTION

Alzheimer's disease (AD) is the most common form of dementia, affecting nearly 38 million people worldwide, a figure that is predicted to double over the next 20 years. The pathological hallmarks of AD are the aberrant deposition of extracellular senile plaques comprised of amyloid- β (A β) peptides and intracellular neurofibrillary tangles (1). Proteolytic cleavage of the amyloid precursor protein generates A β peptides of different lengths (39–43 residues), among which A β_{1-40} and A β_{1-42} are the major isoforms, with A β_{1-42} being more aggregation-prone and toxic than A β_{1-40} (2). The molecular process underlying AD, as outlined in the amyloid β hypothesis, involves an imbalance between production and clearance of A β . This imbalance results in accumulation of amyloid plaques comprised of A β aggregates. The abnormal aggregation of A β peptides into β -sheet-rich fibrils involves a heterogeneous ensemble of oligomeric intermediates, all of which are found to be neurotoxic (3). A β toxicity originates from a number of factors, including formation of ion channels (4), oxidative stress (5), and interaction with receptors (6). NMR studies (7–10) suggest that A β_{1-42} fibrils model as parallel-stacked

hairpin-like structures. The hydrophilic N-terminus (residues 1–16) appears unstructured in those fibril structures, whereas the two hydrophobic patches (residues 17–20 and 30–40) form U-shaped conformations comprised of two intermolecular, parallel, in-register β -sheets separated by a hydrophilic turn region (residues 22–29).

Most proposed pathways for the initial stages of A β amyloid fibril formation suggest some conformational change (misfolding) of the intrinsically disordered A β monomer, resulting in an activated monomer that then recruits other A β molecules to form low-n oligomers (11). Based on the protease-resistant nature of residues A21–A30, this region was proposed as the nucleus site for A β monomer misfolding (activation) (12). A link between monomer misfolding and aggregation propensity and toxicity has been proposed based on experiments (13–16) and simulations (17–26) of the naturally occurring alloforms. Those alloforms contain substitutions that occur mainly within the turn region (residues 22–30). Because of the disordered structure of the N-terminus in the fibril structure, the role of this region in aggregation, toxicity, and pathogenesis was considered for a long time to be negligible (27) and therefore remained largely unexplored. However, several recent reports indicate the possible importance of the N-terminus in A β structure, association, and toxicity (28–30). A novel structural motif

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*Correspondence: daspa@us.ibm.com

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of triple β -sheet within A β_{1-42} amyloid oligomers has been reported with minimal exposure of hydrophobic residues, in which association was found between the N-terminus and residues 17–22 (31). Antibodies that specifically bind to the N-terminus are known to effectively interact with both soluble and insoluble forms of A β (32). Amyloid inhibitor tetrapeptides are also known to bind at the A β N-terminus (33). Two N-terminal mutations, English (H6R) and Tottori (D7N), are reported to alter the monomer misfolding and produce oligomers that are toxic compared with the wild-type (WT) (16). A few recent studies have also suggested the importance of the extreme N-terminus. Experiments reported delayed assembly kinetics and compactness of the D1Y-substituted A $\beta_{40/42}$ peptides (34) and higher toxicity of an A2F variant (35), a consequence of a more hydrophobic N-terminus. It is known that the double mutation D1E/A2V affects A β_{1-40} fibrillation (36). Pyroglutamate-modified A β (A β (pE3-42)) peptides have also emerged as potential key factors in AD pathology due to their abundance in the AD brain, high aggregation propensity, stability, and cellular toxicity (37). A familial A2V mutation has been reported to cause dementia in homozygous carriers, whereas heterozygous carriers were found to be protected (38). The co-incubation of WT and A2V A β_{1-42} peptides was shown to inhibit toxicity in the same study (38). The A2V A β_{1-42} also has been shown to have a very different aggregation pathway involving annular oligomers with higher hydrophobic exposure and stronger toxicity compared to the WT; the A2V peptides more rapidly formed a structured oligomer, as demonstrated by dynamic light-scattering experiments (39).

Recently, a rare A2T variant was found to protect against AD and cognitive decline in the elderly without AD (40). This is thought to represent the first example of an A β variant that confers protection against AD. The prevailing hypothesis for its protective action is that the mutation (A673T in the amyloid precursor protein or A2T in A β), close to the β -secretase cleavage site, lowers A β production, thereby balancing production with clearance. However, the reported lowering in production was a moderate 20% in heterozygous carriers (40). The downstream effects of this mutation could also be vital to its protective behavior (41).

The effects of the A2V and A2T mutations on A β aggregation and toxicity have just started to emerge, and reports appear somewhat controversial. Two recent studies that employed thioflavin T (ThT) fluorescence monitoring suggest very different effects of the A2T mutation on A β_{1-42} aggregation (42,43). Benilova et al. showed no change in A β_{1-42} aggregation upon either A2T or A2V mutation (43). On the other hand, the results from Maloney et al. suggested a lower aggregation propensity of A2T A β_{1-42} when compared to both WT and A2V with similar aggregation profiles (42). In the context of A β_{1-40} , A2T has been shown to aggregate faster (43,44) or similarly (42) compared to

WT. On the other hand, A2V A β_{1-40} exhibited faster aggregation than the WT (38,42,43). The effects of those mutations on aggregation also have been analyzed in combination with the WT peptide. Although 1:1 mixture of WT/A2V A β_{1-42} showed intermediate aggregation compared to the pure solutions in a study by Messa et al. (39), the same mixture was reported by Benilova and colleagues (43) to demonstrate fast aggregation similar to that observed for WT A β_{1-42} . The latter group reported that a 1:1 WT/A2T mixture exhibited no change in aggregation kinetics (43), but Maloney et al. found an intermediate effect (42).

Maloney et al. (42) also compared the toxicity of WT, A2V, and A2T peptides to cultured neurons and found that WT and A2T peptides exhibited similar toxicity, whereas A2V had somewhat lower toxicity. This is to some extent counterintuitive and contradictory to what was reported for the A2V A β_{1-42} by Di Fede et al. (38). Taken together, the effects of A2V and A2T mutations on early aggregation and toxicity are still not fully clear and seem to be very different depending on the isoform (A β_{1-40} versus A β_{1-42}) and experimental conditions. It is also possible that the current experiments are not sensitive enough to capture the effects of those mutations in the more hydrophobic, aggregation-prone, and toxic A β_{1-42} isoform.

Given the protective and causative effects associated with the A2V and A2T substitutions and the established link between monomer misfolding and early aggregation, it is crucial to obtain a comparative view of the conformational landscapes of those two variants and the WT A β_{1-42} full-length monomers. Even with the sophistication of modern techniques, experimental characterization of the monomeric A β peptides remains challenging due to the intrinsically disordered and rapidly interchanging nature of the system. On the other hand, molecular dynamics (MD) simulations at different resolutions have been widely used to complement experiments (45) in providing detailed information on the structure of various A β species ranging from monomers (23,46–50) to oligomers (51,52) to protofibrils (53) to fibrils (54). Interactions of different A β species with toxicity and aggregation inhibitors (55–57) and with lipid bilayers (58,59) also have been studied using MD. In this study, we have performed extensive all-atom replica-exchange molecular dynamics (REMD) simulations of WT, A2V, and A2T A β_{1-42} in explicit water to characterize the monomeric ensemble. To our knowledge, this is the first study comparing the structural landscapes of the A2T and A2V A β_{1-42} monomers. Our simulations reveal an enhanced double-hairpin population in A2V due to hydrophobic clustering between the N-terminus and distant hydrophobic patches. In contrast, the A2T mutation engages the N-terminus in an unusual set of long-range electrostatic interactions with residues such as K16 and E22, resulting in a population of unique conformations that comprise only the C-terminal hairpin. These findings imply that the A2X (X = V or T)

mutations situated within the mainly disordered N-terminus can dramatically alter the WT monomeric conformational landscape by globally rewiring the intramolecular interaction network within $A\beta_{1-42}$.

MATERIALS AND METHODS

Details of the MD simulations regarding system setup, simulation parameters, simulation length, and so forth can be found in the [Supporting Material](#). The total number of replicas used was 64, and each replica was 175 ns long, resulting in an aggregate simulation time of 11.2 μ s per system. A number of convergence checks were performed (see the [Supporting Material](#)).

RESULTS

In this work, we compare *in silico* the monomeric conformational ensemble of the A2V and A2T $A\beta_{1-42}$ variants with that of WT by performing extensive atomistic REMD simulations in explicit water. All three variants exhibit similar secondary structure propensity, with coils and turns being prevalent (see [Fig. S3 a](#) in the [Supporting Material](#)). The secondary-structure propensity values for WT are ~30% coil, ~45% turn, and ~18% β -strand; A2V and A2T show similar values. This estimated composition for the WT system in this study agrees well with an earlier report (20), implying that the sampling obtained using the protocol described here represents the WT monomeric structure

reasonably and consistently well. The ensemble-averaged secondary structure per residue for each variant is shown in [Fig. 1, a–c](#). The residue-based secondary-structure propensity analysis ([Fig. 1, a–c](#)) suggests a primarily disordered N-terminal region (residues 1–15) for all three variants, with a strong turn character around residues 8–9 and 13–15 and a small (5–10%) β -strand propensity in the first 12 residues. Some helical structure near residues 12–16 is also observed. Residues 17–20 (the central hydrophobic cluster (CHC)) and 31–40 show high β -strand propensity ($\geq 25\%$) in addition to a predominant turn around residues 23–30 in the WT ([Fig. 1 a](#)), consistent with the reported strand-loop-strand (SLS) structure of $A\beta_{10-35}$ (60). Interestingly, A2V demonstrates higher β -strand propensity within residues 16–20, 30–35, and 39–41 compared with the WT. A significant increase in the turn propensity compared to the WT also can be noticed in residues 23–30 and 36–37 of A2V ([Fig. 1 b](#)). On the other hand, the β -strand propensity is found to be lower in A2T around residues 18–19, 31–32, and 36–39 ([Fig. 1 c](#)). In addition, the turns around residues 23–30 and 36–37 appear more prominent in A2T. These data indicate that a mutation in the extreme N-terminus (at position 2) can alter the secondary structural preference of the distant hydrophobic regions (the CHC and the C-terminus).

The C_{α} -contact map ([Fig. 1 d, upper triangle](#)) reveals the presence of strong antidiagonal contacts between the CHC and C-terminus, suggesting a β -hairpin formation in the

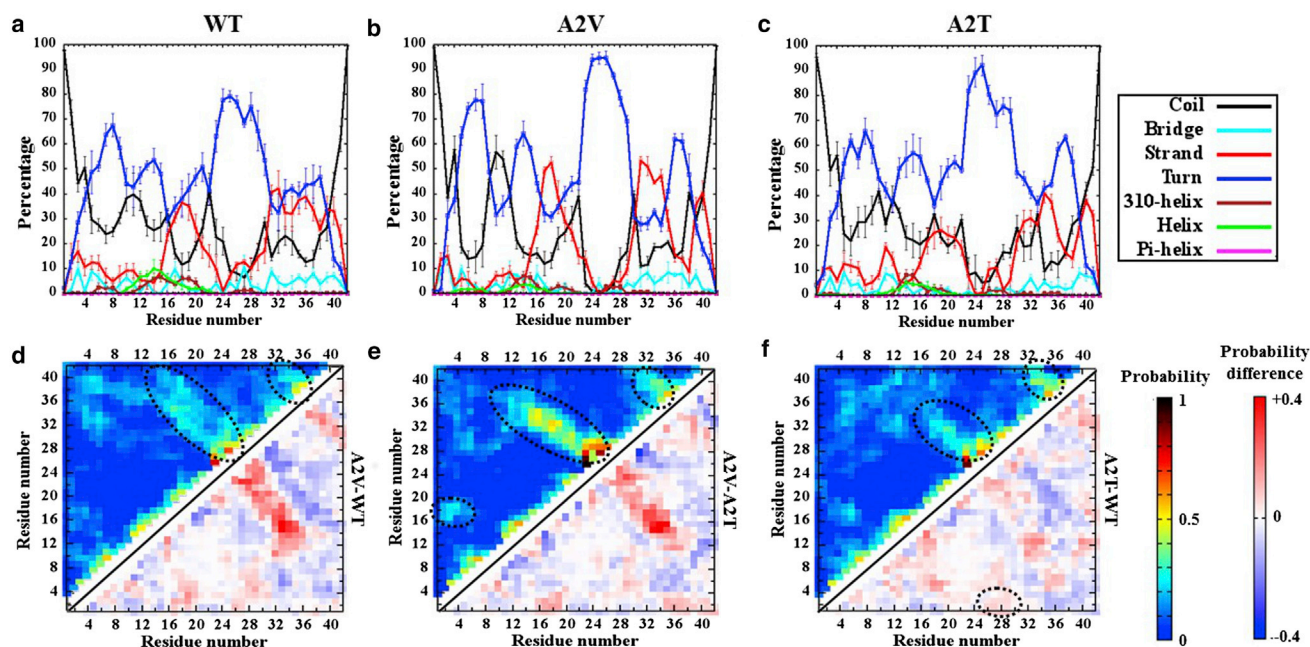


FIGURE 1 Secondary and tertiary structure. (Top) Secondary structure per residue averaged over the production ensemble: (a) WT, (b) A2V, and (c) A2T. Colors used are shown on the right. The standard deviations were estimated by splitting the data in four equal sets. (Bottom) Tertiary contact maps: (d) WT, (e) A2V, and (f) A2T. (Upper triangle) Ensemble-averaged probability of interresidue C_{α} - C_{α} contact formations (nonsequential contacts, i.e. $|i-j| \geq 3$, are only shown) are colored according to the color map shown on the right. (Lower triangle) Arithmetic difference between the contact probabilities for each pair of $A\beta$ variants colored according to the color map shown on the right. Black dotted circles highlight the two sets of antidiagonal contacts as well as the crucial contacts involving residues 1–5, which help discriminating the variants. To see this figure in color, go online.

WT, consistent with earlier simulations (60). β -hairpin formation of residues 17–36 during early A β aggregation has been further supported by an affibody-A β complex structure (61) and by the presence of a turn-like structure at residues 25–29 in A β oligomers (62). Additional antidiagonal contacts appear prevalent in the C-terminal region between residues 33–36 and 39–41, indicative of a second β -hairpin motif. These results imply a double-hairpin topology, in agreement with the simulations performed by Rosenman et al. (20). Long-range interactions between two termini (residues 1–8 and 32–42) are observed in addition to some contacts between the CHC and the extreme N-terminus (residues 1–5). Highly probable local contacts consistently appear in the turn regions.

Fig. 1 e (upper triangle) reveals a more robust nature of those β -hairpin interactions in A2V compared to the WT, suggesting a favored population of double-hairpin structures in A2V. The long-range interactions between the N-terminus (residues 1–4) and the CHC are stronger in A2V (Fig. 2 b, upper triangle) compared with the WT. The difference contact map (Fig. 1 d, lower triangle) further reveals a register shift toward the N-terminus in A2V compared with the WT for both sets of antidiagonal interactions.

In the A2T variant (Fig. 1 f, upper triangle), the antiparallel β -hairpin interactions between the CHC and residues 30–35 are weaker than in the WT, whereas the C-terminal β -hairpin appears more robust. Also, a cluster of contacts between residues 5–7 and the CHC is observed in A2T. Interactions between residues 1–4 and 34–42 appear weaker in A2T than in the WT (Fig. 1 f, lower triangle). However, residues 1–4 of A2T make contact more frequently with the 23–30 turn (Fig. 1 f, lower triangle, dashed circle); in general, those contacts are weak. At least 25% of those contacts occur with a $\geq 10\%$ probability for the A2T variant, compared to 7% probability in the WT and A2V. Taken together, these results suggest that the A2V/T substitution primarily alters 1) the β -hairpin interactions between the CHC and the C-terminus, termed $NC_{\text{CHC-CTR}}$ and 2) the interactions between the N-terminus (Ntr) (residues 1–5) and the CHC, termed $NC_{\text{Ntr-CHC}}$.

In the following, we compare the potential mean force (PMF) plots (Fig. 2) to discriminate between the conformational ensembles of different variants. It should be noted that as all three peptides are intrinsically disordered, it is nontrivial to choose an optimal set of reaction coordinates for defining their free-energy landscapes. Since from the above analysis, the two sets of contacts, namely $NC_{\text{Ntr-CHC}}$

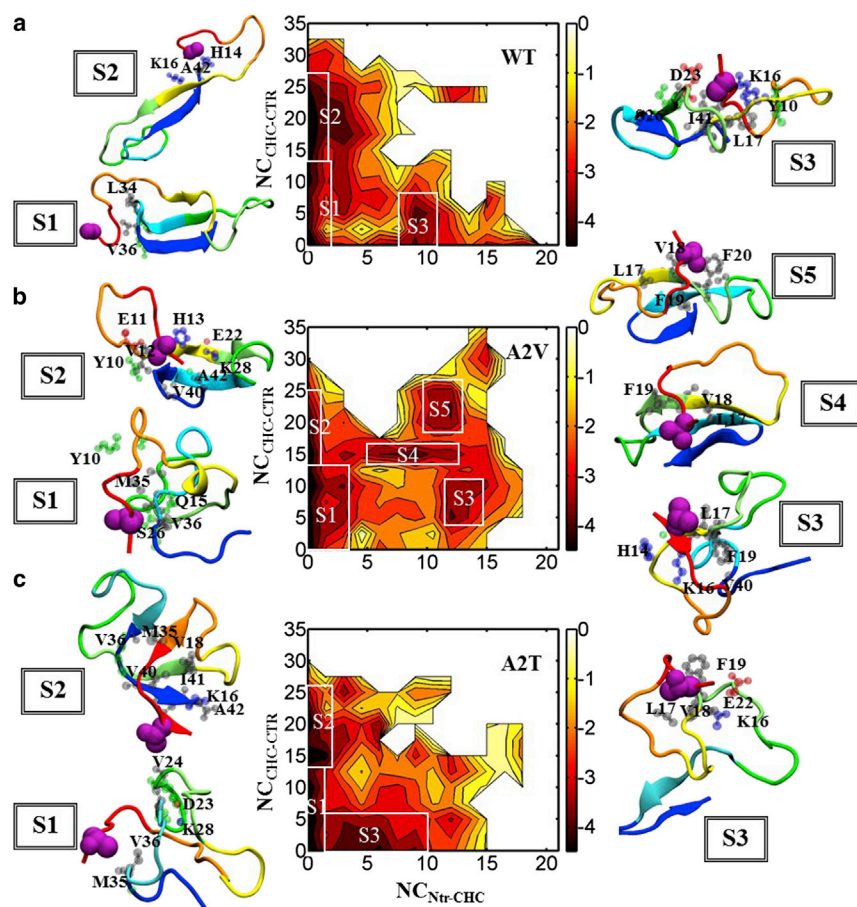


FIGURE 2 Potential of mean force (PMF) plots: (a) WT, (b) A2V, and (c) A2T. Two-dimensional PMF plots as a function of number of contacts between the Ntr (residues 1–5) and the CHC (residues 16–21), $NC_{\text{Ntr-CHC}}$, and number of contacts between the CHC and the C-terminal region (residues 31–42), $NC_{\text{CHC-CTR}}$. Each contour level represents 0.5 kcal/mol. White squares denote the discrete regions on the PMF plots individually representing $\geq 5\%$ of the production ensemble. The representative conformation of the largest cluster for each of those regions is shown using cartoon representation colored with a red-to-blue rainbow spectrum from the N- to C-terminus. Residue 2 is shown in purple spheres, whereas the residues forming long-range contacts with the N-tr are shown using CPK representation. A 5 Å cutoff between heavy atoms is used for this purpose. The residues are colored according to their types (hydrophobic, gray; polar, green; basic, blue; acidic, red). To see this figure in color, go online.

and $NC_{\text{CHC-CTR}}$, appear to capture notable differences in the tertiary contact probabilities among variants, we estimated the PMFs as a function of them (see the [Supporting Material](#)). The WT PMF plot reveals distinct features of the most populated regions (regions S1–S3) ([Fig. 2 a](#), *white boxes*) of the landscape. The S1 region corresponds to the structures with strongly limited $NC_{\text{Ntr-CHC}}$ and $NC_{\text{CHC-CTR}}$ contacts. Structures with robust CHC-CTR interaction accompanied by a few (≤ 5) Ntr-CHC contacts populate the S2 region. In contrast, S3 conformations exhibit stronger Ntr-CHC interaction but lack substantial CHC-CTR contacts, and they are much less populated (compared to S1 and S2). We also performed a cluster analysis with a 3 Å $C\alpha$ -RMSD cutoff distance on those discrete regions. The definition criteria, population size, and cluster analysis results for those regions are listed in [Table S1](#), demonstrating that those structures all together represent $>75\%$ of the total ensemble for each system. The total number of clusters and the size of the top five clusters show that the S1 structures are more disordered than S2 structures, whereas S3 can be well represented by a single cluster. The representative structure for the largest cluster of each region is shown in [Fig. 2 a](#). Additional representative structures, as well as the tertiary contact maps and the residue-based secondary-structure propensities, can be found in

[Fig. S4](#). Those data confirm that the WT S1 conformations are mostly disordered and show very little secondary and tertiary structure ([Fig. S4 a](#)). The only structured region in the S1 conformations is the C-terminus, with a 20% β -hairpin population. In contrast, both sets of hairpin contacts are evident in S2 structures, the C-terminal hairpin being slightly weaker than the CHC-CTR one ([Fig. S4 b](#)). Interestingly, the minor S3 population consists of interaction of the 23–30 turn with both termini, which is absent in S1 and S2 structures ([Fig. S4 c](#)). Representative conformations illustrated in [Figs. 2 a](#) and [S4](#) support these results.

The S1 and S2 regions in A2V ([Figs. 2 b](#) and [S5, a and b](#)) and A2T ([Figs. 2 c](#) and [S6, a and b](#)) monomers show similar features overall when compared to the WT. Interestingly, A2V samples additional structures containing significant Ntr-CHC and CHC-CTR interactions ([Fig. 2 b](#), S4 and S5 regions); such conformations are rarely observed for the WT and A2T. The representative structures from those regions imply that a nonpolar amino acid at position 2 (i.e., Val) forms a hydrophobic cluster involving hydrophobic side chains from CHC, resulting in further stabilization of CHC-CTR contacts ([Fig. 2 b](#)). Accordingly, the secondary-structure analyses and tertiary contact maps for S4 and S5 structures ([Fig. 3](#), *middle and right*) reveal

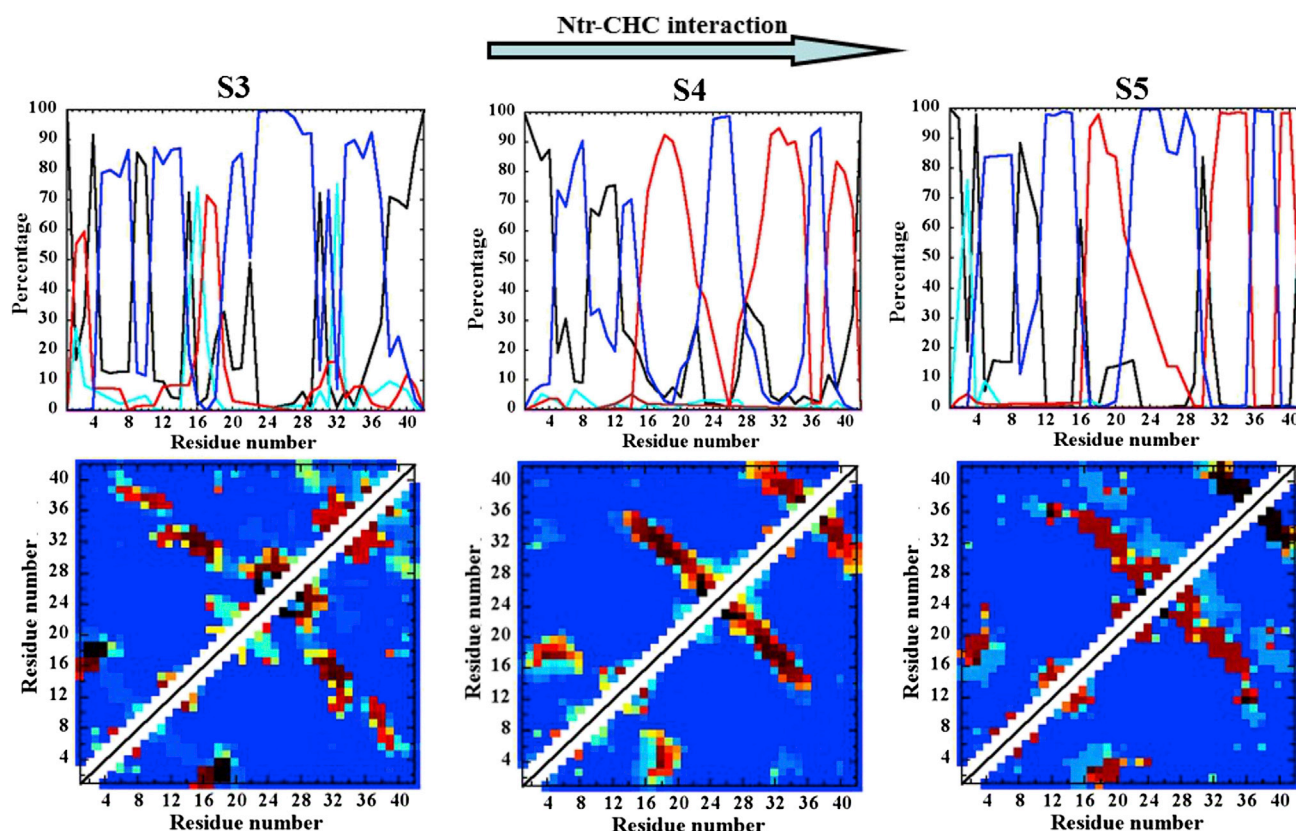


FIGURE 3 Analysis of the A2V conformations with enhanced Ntr-CHC interaction. The residue-based secondary structure and tertiary $C\alpha$ - $C\alpha$ contact maps for A2V conformations corresponding to the S3 (*left*), S4 (*middle*), and S5 (*right*) regions. S4 and S5 structures demonstrate double-hairpin topology triggered by Ntr-CHC contacts. Color schemes used are the same as in [Figs. 1](#) and [2](#). The top arrow indicates the relative probability of Ntr-CHC interaction. To see this figure in color, go online.

double-hairpin-like conformations. Those conformations are regularly populated in A2V (S4 + S5 represent 15% of the total ensemble (Table S1)), as compared with the WT. Lowering of disorder can be seen in those double-hairpin conformations, as revealed from the total number of clusters and the size of the top five clusters (Table S1). Fig. 3 further shows that both sets of hairpin interactions experience a subtle register shift toward the N-terminus in a subpopulation of those double-hairpin structures (the S4 region). The Ntr-CHC contacts are also found in the S3 structures of A2V; however, in those structures, the C-terminus remains in a collapsed coil structure (Fig. 3, left).

In sharp contrast, A2T more frequently visits conformations with significant Ntr-CHC interaction but much weaker CHC-CTR contacts (Fig. 2 c, S3 region). Interestingly, this region of the landscape is rarely populated for both WT and A2V, but the population for A2T is a significant 17% of the total ensemble (Table S1). Further analysis reveals a weaker CHC-CTR interaction resulting in the loss of the first hairpin present in those unique S3 structures (Figs. 2 c and 4, a and b). Consequently, those structures contain only the C-terminal β -hairpin (Fig. 4, a and b). The side-chain-side-chain contact map illustrates strong interaction between the N-terminus (residues 1–6) and residues 16–22 (Fig. S7); however, residue T2 appears to be weakly involved in those interactions and remains solvent-exposed. In particular, strong D1-K16 and R5-E22 side-chain-side-chain contacts are observed. Accordingly, the representative structure illustrates an ionic network comprised of residues D1, E3, R5, K16, and E22 (Fig. 4 c). The salt-bridge population analysis (Table S2) indicates that such ionic interaction becomes more probable in the presence of a solvent-exposed threonine at position 2. For example, the D1-K16 salt bridge is populated with a probability of 34% in S3 structures. At the same time, the formation of the E22-K28 salt bridge becomes less probable (5% (Table S2)) in S3 structures compared to what is observed in S1 and S2 (35% and 12% probabilities, respectively). Taken together, the altered ionic network between the N-terminus and distant charged residues, such as K16 and E22, appears to hinder the formation of the first hairpin interaction in the S3 structures of A2T.

DISCUSSION

Given the reported protective and causative effects associated with A2V and A2T substitutions vis à vis AD, it is of importance to understand the differences in the self-assembly process of those variant peptides, which is intimately associated with the disease pathogenesis. The first step toward this goal is to characterize the conformational landscapes of those variant monomers and compare them with the WT. To our knowledge, this is the first study to compare the structural landscapes of the A2T and A2V A β_{1-42} monomers. Our findings from extensive REMD simulations

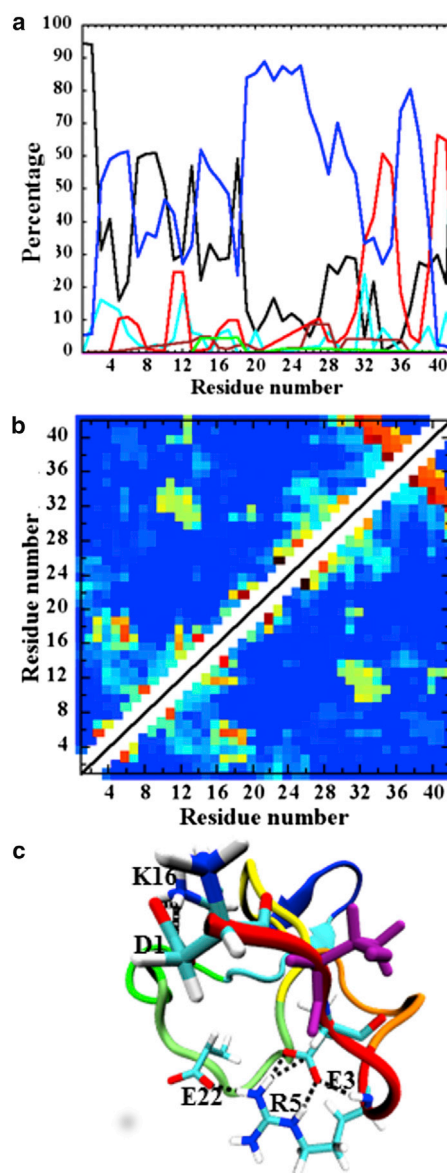


FIGURE 4 Unique A2T conformations with altered ionic network. (a) Residue-based secondary structure, (b) tertiary Ca-Ca contact map, and (c) ionic network within the representative conformation of the largest S3 cluster. The residues, i.e., D1, E3, R5, K16, and E22, forming long-range ionic interaction, are shown using licorice representation. Hydrogen bonds are shown in black dotted lines. Residue Thr2 is shown in purple. Color schemes used are the same as in Figs. 1 and 2. To see this figure in color, go online.

report significant differences in terms of the transient structure population in those variants, which is modulated by interactions of the N-terminus with hydrophobic residues (the CHC) in the case of A2V or with charged residues (K16 and E22) in the case of A2T.

The more persistent β -hairpin character in the hydrophobic regions that lead to a double-hairpin topology in the A2V monomer, as revealed in our simulations, could have important implications in determining the aggregation and

toxicity of A2V. This monomer has been reported to follow a peculiar aggregation behavior that leads to the formation of annular structures with a higher hydrophobicity profile and hence higher toxicity *in vitro* (39) as well as *in vivo* (63). A2V also showed greater β -conformation propensity during aggregation (64). The formation of β -structures is believed to be important in amyloid aggregation and toxicity. For example, β -hairpin monomeric conformations in amyloid peptides are found to represent metastable states that interconvert slowly and thus can act as the seed for nucleation (65). Experiments have also highlighted the importance of β -hairpin stabilization in the A β monomer in inducing a toxic soluble oligomer population (61,66,67). NMR and atomic force microscopy (AFM) experiments (68) have suggested the presence of double-hairpin conformations within toxic A β_{1-42} oligomers. Double-cysteine mutants (A $\beta_{40_{cc}}$ and A $\beta_{42_{cc}}$) with overly stable β -hairpin monomeric conformation have been shown to lower fibril formation and increase populations of toxic β -sheet oligomers and/or protofibrils (67). A structural comparison of monomeric A β_{1-42} and fibrillar A β_{1-42} suggests that the monomeric hairpin conformation needs to retain sufficient flexibility to undergo a concerted conformational change toward the extended parallel β -sheet structure found in fibrils (61), as also observed in MD simulations of A β_{10-35} dimers and trimers (66). The fact that double-hairpin structures were only observed in A β_{1-42} and not in A β_{1-40} has been linked to the higher aggregation propensity and toxicity of the longer isoform (20). Taken together, our simulations, combined with results of previous studies, imply that the transiently formed double-hairpin structures of the A2V monomer might contribute to the stabilization of small, toxic oligomers composed of β -structures, thus inhibiting fibrillation. Thus, a single-point nonpolar substitution at position 2 of A β_{1-42} might act as a switch to redirect the peptides to aggregate via a neurotoxic pathway by stabilizing the β -hairpin conformation (Fig. 5). Our results are consistent with simulations of the shorter A2V A β_{1-28} monomer, which indicated that a valine at position 2 can favor Ntr-CHC interaction (69). Nevertheless, the simulations of the full-length A2V peptide presented here demonstrate how the hydrophobic Ntr-CHC interaction can lead to enhanced CHC-CTR hairpin population, leading to the double-hairpin topology (Fig. 5).

Our simulations further suggest a competition between specific hydrophobic and electrostatic interactions modulated by the N-terminus within the A β_{1-42} monomer, which in turn regulates the delicate balance among the local minima present in its frustrated conformational landscape (Fig. 5). As discussed above, the N-terminus in the A2V variant stabilizes the double-hairpin conformation by clustering with distant hydrophobic residues (Fig. 5). In contrast, the A2T variant, with its more polar N-terminus, leads to unique structure populations that are rarely seen for the WT and A2V. Although Thr-2 itself minimally inter-

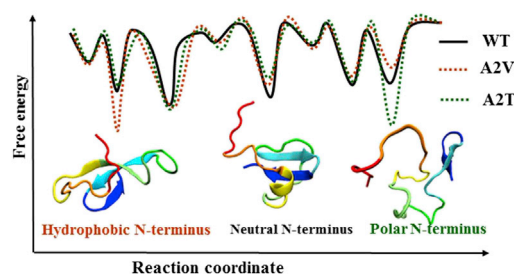


FIGURE 5 Schematic illustration of the A β_{1-42} N-terminus influence on the monomeric landscape. The curves show free energy of the variant monomers in one dimension. Based on the analysis, the reaction coordinate might represent a combination of the CHC-CTR interaction and Ntr-peptide interaction. The N-terminal substitution, depending on its hydrophobic or polar nature, can switch the relative population among different β -hairpin conformational states present in the WT landscape. It should be noted that this is an overly simplified representation of the A β_{1-42} landscape, and many other conformational states are present due to the intrinsically disordered nature of the system. To see this figure in color, go online.

acts with the rest of the protein in those conformations, its presence allows the N-terminus (e.g., residues D1, E3, and R5) to form an unusual set of ionic interactions with distant residues, such as K16 and E22. This effect impairs the formation of the E22-K28 salt-bridge that is known to be crucial for CHC-CTR hairpin formation (18). In consequence, those structures comprise only the C-terminal short β -hairpin (Fig. 5).

Mutations at residues E22 (E22K, E22G, and E22Q) and D23 (D23N) that affect A β aggregation and cause early-onset AD (15) are reported to alter the WT 21–30 turn structure in simulations (70). Two N-terminal mutations (H6R and D7N) also have been found to affect the extent of β -structure and salt-bridge network within the monomer (71,72). Nevertheless, those mutations directly change the charge distribution within the peptide, which is not the case for A2T. At first glance, a threonine at position 2 would not be expected to have a dramatic effect on the A β_{1-42} conformational landscape by altering the electrostatic interaction network. It appears that Thr-2 changes the local conformational preference of the N-terminus, consequently promoting alternative electrostatic interactions that compete with those required for turn nucleation and CHC-CTR hairpin formation. This leads to population of rare and unique conformations (S3 structures). Since β -hairpin abundance has been linked with aggregation mechanisms and toxicity associated with AD (17,73), the unique A2T conformations devoid of the CHC-CTR β -hairpin that also have an altered electrostatic interaction network might contribute toward its slower aggregation (42). It is noteworthy that a threonine in position 2 appears to introduce more disorder within A β_{1-42} , whereas a valine at the same position induces some order, which overall is consistent with the intrinsic (dis)order-promoting nature of those amino acids (74). However, our simulations reveal that this effect arises from the global interaction changes triggered by a single

amino acid substitution. Interestingly, conformation-specific antibody staining experiments imply the involvement of the N-terminus in tertiary interactions in the oligomers of both A2V and A2T variants (43). Our simulations demonstrate that such an interaction is also present at the monomer level and is strikingly different between the two variants, which might play a critical role in their early aggregation by populating different monomer structures. Since the A β N-terminus also has been suggested to be crucial in binding with membrane (75) and with some toxic A β receptors (76), the altered interaction pattern observed in the A2V and A2T monomers may also affect their toxicity (77).

If the interaction between the extreme N-terminus and the rest of the peptide is indeed critical for aggregation and toxicity, then it may be possible, for example, to use small organic molecules (78) and peptides (79,80) that would bind to A β monomers and/or oligomers and inhibit aggregation and/or toxicity by intervening in or altering this interaction. The role of the extreme N-terminus in modulating the A β_{1-42} monomeric hairpin structure revealed here may guide the design of inhibitory peptides, including variants of the WT (¹DAEFRH⁶) sequence (64).

CONCLUSIONS

Taken together, our findings suggest that the effect of the second amino acid on the A β_{1-42} monomer structure is highly complex and sequence-dependent. An enhanced double-hairpin population similar to those reported in toxic WT A β_{1-42} oligomers is found in the A2V monomer. Hydrophobic clustering between the N-terminus and the central and C-terminus hydrophobic patches promotes such double-hairpin formation in A2V. In contrast, the A2T mutation triggers unusual ionic interactions of the N-terminus with K16 and E22, thereby impeding CHC-CTR hairpin formation. Consequently, a unique population comprising only the C-terminal hairpin is observed. Although further investigation is needed to obtain a complete molecular picture of the relationship between monomer misfolding, aggregation, and toxicity, and protection against or causation of AD by these N-terminal variants, the simulations described here clearly show that single A2V and A2T substitutions can alter the structural landscape of the A β_{1-42} monomer by shifting the equilibrium to different conformational states.

SUPPORTING MATERIAL

Supporting Material, seven figures, and two tables are available at [http://www.biophysj.org/biophysj/supplemental/S0006-3495\(14\)04758-4](http://www.biophysj.org/biophysj/supplemental/S0006-3495(14)04758-4).

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