

Structure and Dynamics of Parallel β -Sheets, Hydrophobic Core, and Loops in Alzheimer's A β Fibrils

Nicolae-Viorel Buchete and Gerhard Hummer

Laboratory of Chemical Physics, National Institute of Diabetes and Digestive and Kidney Diseases, National Institutes of Health, Bethesda, Maryland 20892-0520

ABSTRACT We explore the relative contributions of different structural elements to the stability of A β fibrils by molecular-dynamics simulations performed over a broad range of temperatures (298 K to 398 K). Our fibril structures are based on solid-state nuclear magnetic resonance experiments of A β (1–40) peptides, with sheets of parallel β -strands connected by loops and stabilized by interior salt bridges. We consider models with different interpeptide interfaces, and different staggering of the N- and C-terminal β -strands along the fibril axis. Multiple 10–20 ns molecular-dynamics simulations show that fibril segments with 12 peptides are stable at ambient temperature. The different models converge toward an interdigitated side-chain packing, and present water channels solvating the interior D23/K28 salt bridges. At elevated temperatures, we observe the early phases of fibril dissociation as a loss of order in the hydrophilic loops connecting the two β -strands, and in the solvent-exposed N-terminal β -sheets. As the most dramatic structural change, we observe collective sliding of the N- and C-terminal β -sheets on top of each other. The interior C-terminal β -sheets in the hydrophobic core remain largely intact, indicating that their formation and stability is crucial to the dissociation/elongation and stability of A β fibrils.

INTRODUCTION

Fibrils of Alzheimer's amyloid-beta (A β) peptides are the major component of Alzheimer's disease plaques (1,2). Amyloid fibrils are also associated with other neurodegenerative and prion diseases, type 2 diabetes (2–4), and insulin deficiency in the brain (5). In Alzheimer's disease, recent studies suggest a connection between A β plaques, oligomers, and neurotoxic effects (6), but its exact nature remains largely unknown (7). The detailed structural characterization of A β peptides both in solution (8–10) and in fibrils is a crucial step toward understanding the formation and stability of ordered fibrillar peptide aggregates (2,11–13). Knowledge of structural details of A β fibrils should in particular facilitate the design of inhibitors of fibril formation (14–16). Moreover, insights into the molecular self-assembly processes during fibril formation may also aid in the development of new soft nanomaterials (17,18).

Most computational investigations of the structure and dynamics of A β peptides have focused on monomers (10,19), dimers (20–22), and other low-order oligomers (20,23–25). By studying early aggregation events, molecular simulations using simplified, coarse models of various amyloid-forming molecules provided detailed mechanistic and structural insights in the formation of prefibrillar amyloid species (21, 26–29). Larger systems have been studied to identify general aspects of amyloid assembly (30–35).

Here, taking advantage of structural information available from recent solid-state nuclear magnetic resonance (ssNMR) experiments (11,13,36,37) on the parallel cross- β structure

of A β _{1–40} protofilaments, we perform all-atom/explicit solvent simulations of amyloid fibrils containing six two-peptide units, each with two U-shaped A β _{9–40} peptides in a plane roughly perpendicular to the fibril axis. Among the different structural topologies observed depending on experimental growth conditions (11,13,37), we focus on A β _{1–40} fibrils grown under gentle agitation. In a series of different models, we incorporate information from recent isotope-dilution ssNMR (11) suggesting that the N- and C-terminal β -strands within a given peptide may not be in contact, unlike previous models.

The objectives of our study are twofold: 1), to explore the contributions of the different structural elements of typical A β protofilaments to stability, conformational dynamics, and elongation/dissociation mechanisms; and 2), to investigate different possible protofilament models, including different β -sheet staggering and loop conformations, searching for common features that may be independent of structural details of specific models. Our simulations of both wild-type and mutated sequences cover a broad temperature range (298, 348, and 398 K), in which fibrils are experimentally stable (below \sim 330 K) or fully dissociate (above \sim 373 K) (38,39). We discuss structurally relevant fibril characteristics such as the secondary-to-quaternary structural elements (e.g., β -strands, intra- and intermolecular contacts), internal salt-bridges, the conformations of the amino acids in the loop region, and the interior hydration. We analyze the evolution of these elements at elevated temperatures, focusing in particular on the structure and dynamics of the A β monomers at the fibril ends to identify possible dissociation/elongation mechanisms.

The outline of the article is as follows. After introducing fibril terminology, we describe the different structural

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Address reprint requests to G. Hummer, Tel.: 301-402-6290; E-mail: hummer@helix.nih.gov or gerhard.hummer@nih.gov.

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models and the simulation methodology. We then present simulation results for infinite periodic fibrils. From simulations of fully solvated finite fibrils, we extract information about secondary structure, side-chain packing, and interior salt bridges and hydration as well as fibril dynamics. We then interpret our results obtained at different temperatures in the context of fibril stability and elongation/dissociation mechanisms.

METHODS

In studies of A β amyloid aggregates, the term “fibril” is generally used for structures of aggregated peptides that have a rodlike appearance and low or unknown degree of molecular organization (12,40). The term “protofilament” refers more specifically to basic structural units of amyloid fibrils, consisting of A β peptide monomers arranged in two molecular layers along the fibril axis (12). Here, we study the molecular structure of A β peptides in such protofilaments and use the terms “fibril” and “protofilament” interchangeably. “Two-peptide” units refer to the two A β_{9-40} peptides in a plane roughly perpendicular to the fibril axis.

Structural models of A β protofilaments

We use A β_{9-40} to model the full-length A β_{1-40} protofilaments (Fig. 1 A). N-terminal residues 1–8 are structurally disordered and not necessary for fibril growth (12,36,37). Structural models of A β protofilaments were built using peptide structures taken from the simulations of Buchete et al. (12) that were then stacked along the fibril axis (Fig. 2).

We explore different staggers (11,12,35) of the N- and C-terminal β -strands along the fibril axis to account for recent isotope-dilution experiments (11). We study three systems (S1, S2, and S3) that differ in the

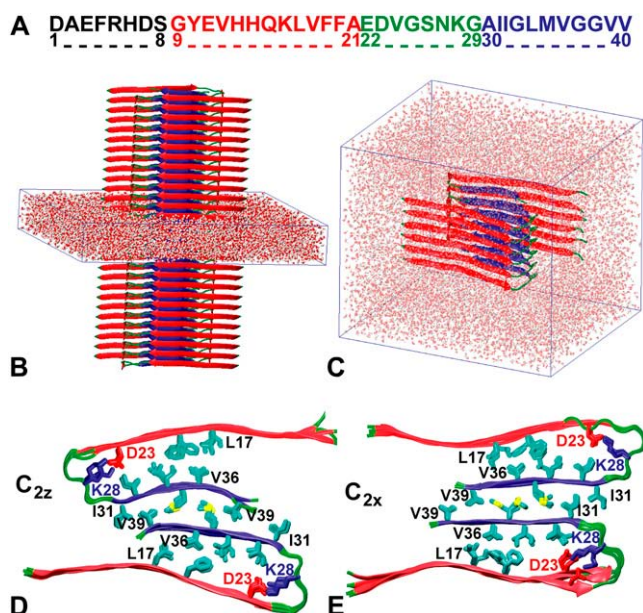


FIGURE 1 MD simulations of A β_{9-40} protofilaments. (A) A β_{1-40} sequence and major structural elements: the unstructured N-terminal region (black), the N- (red) and C-terminal β -strands (blue), and the loop region (green). (B) Infinite-periodic fibril with solvent-filled simulation box. (C) Solvated 12-peptide fibril segment. (D,E) Top views of infinite fibrils with C_{2z} (D) and C_{2x} (E) topologies after 10 ns of MD.

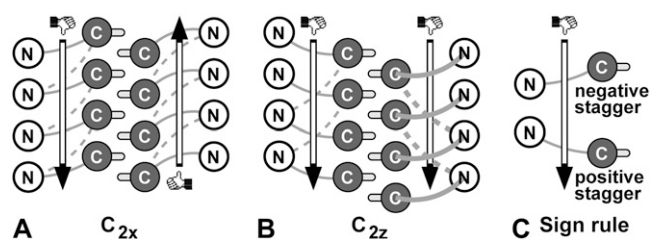


FIGURE 2 Schematic of A β_{9-40} protofilament β -strand staggering of fibrils with (A) C_{2x} and (B) C_{2z} topologies. The N- (open) and C-terminal β -strands (dark shaded circles) are roughly perpendicular to the plane. Thick and thin shaded connecting loops are located in front and behind, respectively. Solid loops correspond to the initial internal staggering of the S1 models (−0.5 for S1-C_{2x} and +0.5 for S1-C_{2z}), and dashed loops indicate −1.5 staggering of S2 and S3. M35 side chains (light shaded) indicate the 0.5 external staggering. Vertical arrows indicate the direction of the two fibril strands, defined by a right-hand rule for the N-to-C direction of the A β peptide.

initial staggering of their N- and C-terminal β -strands (+0.5 for S1 and −1.5 for S2 and S3). Staggering is defined as the β -strand displacement along the oriented fibril axis in units of the ~ 4.8 Å β -sheet interstrand spacing (11) with the sign determined by the intrinsic fibril direction (Fig. 2 C). A right-hand rule for the N-to-C direction of the peptide (Fig. 2 C) is used to define the staggering sign and the intrinsic fibril direction. Note that the staggering signs obtained with this rule are opposite to the ones depicted in Fig. 7 of Petkova et al. (11). To change the initial staggering from +0.5 (S1 system) to −1.5 (S2 and S3 systems), we reconnect the V24–N27 loops of stacked β -strands (Fig. 2). The S2 and S3 systems have different structures of the connecting loop, with V24–N27 being more extended in the S3 system. Additional simulations are performed for an S2 system with mutations V24A, S26A, and N27A.

For each A β fibril model, 10 ns of molecular dynamics (MD) simulations at 298 K were conducted in an infinite-periodic fibril setup (Fig. 1 B), followed by multiple simulations of finite fibril segments (Fig. 1 C) at 298, 348, and 398 K for up to 20 ns.

For S1 and S2 systems, we simulate both C_{2x} and C_{2z} topologies, which differ in the relative orientations of the A β peptides in a two-peptide unit normal to the fibril axis (see (12), and Fig. 1, D and E). For S3, we only consider the C_{2z} topology that is more strongly supported by experiment (11). The two topologies differ in the relative orientation of the contacting C-terminal β -strands in the interface between the two U-shaped peptides within a two-peptide unit: the two strands are parallel in C_{2x} structures, and antiparallel in C_{2z} structures. As a consequence, the two stacks of U-shaped peptides along the fibril have either parallel (C_{2z}) or antiparallel orientation (C_{2x}), with equivalent fibril ends in the case of C_{2x} fibrils (Fig. 2 A), but differing end structures in the case of C_{2z} fibrils (Fig. 2 B). However, if the staggering sign or magnitude (Fig. 2 C) differ in the two molecular layers of the protofilament (as seen in some of our high-T simulations), the two fibril ends can differ even in the C_{2x} case. We note that differences in the structure of the two ends may affect the relative fibril elongation kinetics. Similarly, symmetry differences between C_{2x} and C_{2z} topologies result in having the structurally disordered N-terminal residues (12,37) either close to each other (C_{2x}, Fig. 1 E) or on the opposite sides of A β protofilaments (C_{2z}, Fig. 1 D), with possible implications on their lateral aggregation properties.

Molecular dynamics simulations

Our molecular dynamics (MD) simulations of A β amyloid protofilaments use the methods described in Buchete et al. (12). Here, we study systems of 12 A β_{9-40} peptides (as compared to eight peptides in (12)), at both ambient (298 K) and elevated temperatures (348 K and 398 K). All simulations of solvated amyloid fibrils were performed using the NAMD2 program (41)

with the CHARMM27 (42) force field parameters. The fibrils were explicitly solvated with TIP3P water molecules (43). All simulations were performed in the NPT ensemble. The Langevin piston method (41,44,45) was used to maintain a constant pressure of 1 atm. The temperature was controlled by using Langevin dynamics with a coupling coefficient of 1 ps (41). We used periodic boundary conditions and the particle-mesh Ewald method (46) with a real-space cutoff distance of 10 Å and a grid width smaller than 1 Å. The switching distance for nonbonded electrostatics and van der Waals interactions was 8.5 Å with a cutoff distance of 10 Å, and the integration time step was 1 fs (with the exception of the S3- C_{2x} system where 2 fs time steps were used in conjunction with constrained bonds of hydrogen atoms (47)).

Initially, between 10 and 20 ns of MD were performed for four-layer infinite-periodic fibril systems hydrated with up to 7071 water molecules. From those systems, six-layer fibril segments were extracted and fully solvated with up to 21,913 water molecules leading to simulated systems of ~70,000 atoms. After successive stages of minimization, heating and equilibration, up to 20 ns of MD runs were performed at 298 K. Additional runs of 10–20 ns were performed at 348 K and 398 K. The combined simulation time for all three systems (S1, S2, and S3) and temperatures is 320 ns (i.e., 80 ns for the infinitely-long systems and 240 ns for simulations of protofilament segments containing 12 $A\beta_{9-40}$ peptide monomers).

RESULTS AND DISCUSSION

β -sheet staggering in infinite-periodic fibrils

After 10 ns of MD, the infinite-periodic fibrils S1, S2, and S3 maintain their secondary structure and overall topology despite a compaction in the side-chain packing (Fig. 1, *D* and *E*, and Fig. S1 that is published as Supplementary Material). In all cases, the side chains of the N- and C-terminal β -strands interdigitate during the equilibration, resulting in a well-packed structure. During the initial equilibration at ~100 ps in the 298 K simulation of the infinite system with S1- C_{2x} topology, the staggering changed from +0.5 to -0.5, indicating a cooperative sliding of both N-terminal β sheets relative to the two C-terminal sheets.

T-dependent secondary structure

From the final structure of the infinite-periodic MD runs, stacks of six two-peptide units (i.e., 12 $A\beta_{9-40}$ monomers) are separated and fully solvated. In subsequent NPT MD simulations, the resulting finite protofilament fragments are free to twist about the fibril axis, unlike the infinite-periodic fibrils. Fig. 3, *A* and *B*, illustrate the time evolution of the secondary structure content for the $A\beta$ segments of the S1- C_{2x} system along the 10 ns trajectory at 298 K and at 398 K, respectively.

For all three models, S1, S2, and S3, both N- and C-terminal β -sheet regions are stable at room temperature for the duration of the simulations (Fig. 3 *A*, and Figs. S2 *A* and S3 *A*). Even peptides located at the ends of six-layer fibril segments present stable β -strand regions. Occasional loss of β -sheet content is more common in the S2 and S3 systems, especially in the loop regions.

However, at elevated temperatures (Fig. 3 *B*, and Figs. S2 *B* and S3 *B*) there is a significant loss of β -sheet content in all cases. Consistent with the interpretation of calorimetry

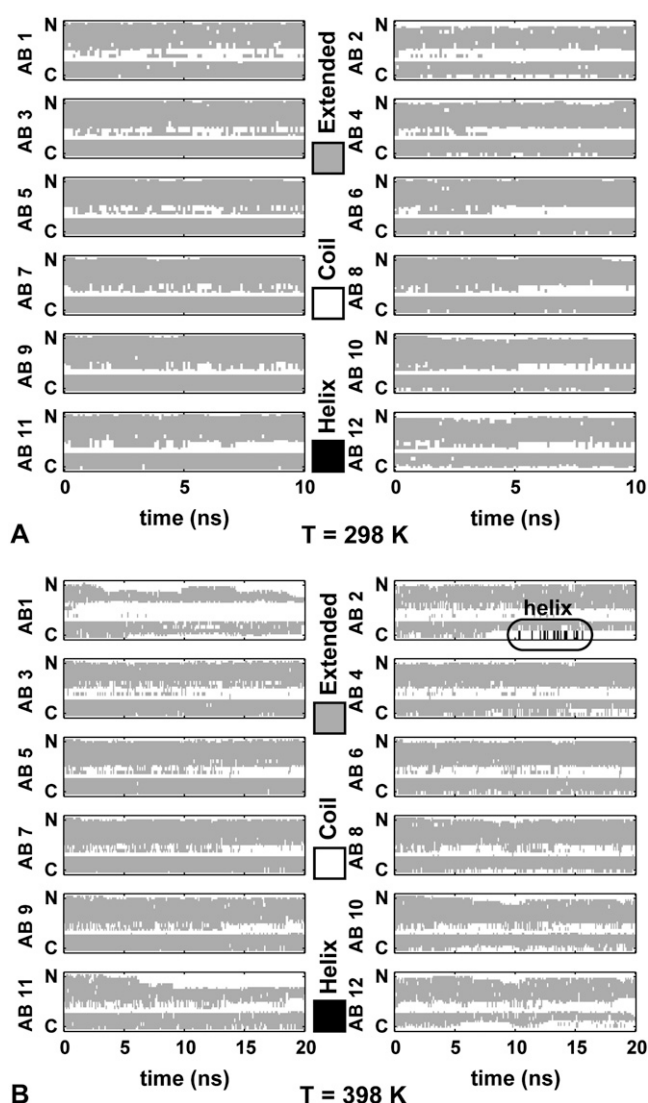


FIGURE 3 Secondary structure (60) of the S1- C_{2x} system at (A) 298 K and (B) 398 K. Shaded representation indicates extended β -strand regions, and open representation indicates coil. Helical segments shown in solid representation are boxed. Peptides AB1/AB2 and AB11/AB12 form the top and bottom layers of the protofilaments, respectively.

experiments (38,39), temperatures above 100°C are sufficient to overcome the hydrophobic interactions between the β -sheet residues, eventually dissociating the fibrils. The interior of the S1- C_{2x} system, starting with +0.5 staggering, presents smaller changes than the S2 and S3 systems initiated with -1.5 staggering. Snapshots of initial and final conformations in MD trajectories of up to 20 ns, illustrating the structural elements of the $A\beta$ protofilament systems, are shown in Figs. S4–S8.

“Surface melting” of N-terminal β -strands

At elevated temperatures, the loss of structure is initiated at the surface of the protofilaments (Fig. 3 *B*), as seen also in

proteins (48). The solvent-exposed N-terminal β -strands are found to be most susceptible to such “surface melting.” In contrast, the interior C-terminal strands maintain their structure for up to 20 ns at all temperatures. Even at the fibril ends, the C-terminal strands appear more stable—an observation with possible implications on the potential fibril dissociation/elongation mechanism, as discussed below in more detail.

External and internal staggering of β -strands in A β fibril segments

For all systems studied here, the C-terminal β -strands of the two interior β -sheets are displaced along the fibril axis by ~ 2.4 Å, corresponding to an external staggering of 0.5. The odd-numbered side chains I31, M35, and V39 adopt an interdigitated packing pattern along both the fibril axis and the β -strand direction. MD runs at elevated temperatures show that the C-terminal strands of different peptides form a highly stable protofilament core.

The internal staggering between the N and C-terminal β -strands of a peptide remains unchanged at 298 K (+0.5 for S1-C_{2z}, -0.5 for S1-C_{2x}, and -1.5 for S2 and S3). However, at elevated temperatures, we observe a tendency of the S2 and S3 systems to go from -1.5 to +0.5 staggering.

The most dramatic change in the staggering of N- and C-terminal strands occurs at 398 K in the S1-C_{2z} system. Starting from +0.5 staggering, a whole N-terminal sheet is collectively displaced after ~ 5 ns, and then again at 8 and 14 ns. As the N-terminal sheet slides on top of the C-terminal sheet, the staggering of the β -strands changes from +0.5 to +1.5, then +3.5, and back to +2.5. The concerted β -sheet sliding is evident in the time-dependent Euclidian distances $d_{i,i+k}$ between terminal heavy atoms of facing side chains of N- and C-terminal β -strands of molecules i and $i+k$ (Fig. 4). In our simulations, we note that the staggering changes less for residues close to the loop than for those at the tip of the strands (Fig. 4D), with N- and C-terminal strands being at an angle to each other.

D23/K28 salt bridge

Consistent with the ssNMR data (11,13,36), all our starting fibril conformations have D23/K28 salt bridge contacts in the loop region. Other studies also showed that charged residues are important for the dynamics of protein aggregation and the stability of β -sheet structures (32,49–51). Consistent with previous results (12), our trajectories of both infinite and finite fibril segments show that the D23/K28 bridges are maintained at 298 K. Positive and negative charges alternate along the fibril axis as in a one-dimensional ionic crystal, with +0.5 staggering for the S1 system, and -1.5 for S2 and S3. We note that D23/K28 salt bridges maintain their staggering even in high-T simulations where the N- and C-terminal β -sheets slide on top of each other.

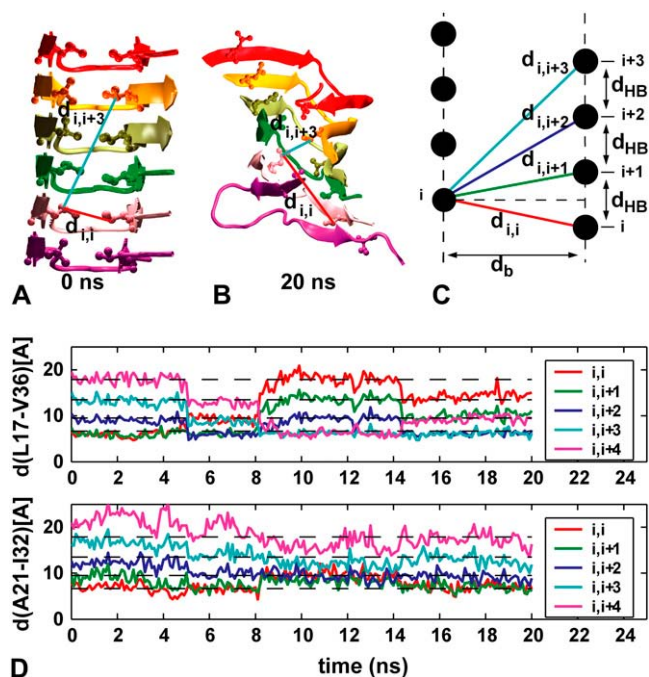


FIGURE 4 Sliding of N- and C-terminal β -sheets in S1-C_{2z} system at 398 K. (A) Starting configuration. (B) Configuration after 20 ns. (C) Schematic of $d_{i,i+k}$ distances. (D) $d_{i,i+k}$ as a function of time for residue pairs L17-V36 and A21-I32 far from the loop and close to it, respectively. Dashed lines indicate $d_{i,i+k}$ distances expected for ideal β -sheets.

Water channels along fibril axis

In the infinite-periodic fibrils, small cavities form near the D23/K28 salt bridges but do not fill with water. However, water penetrates into finite fibril segments to form narrow water channels solvating the interior D23/K28 salt bridges. Indirect experimental evidence for interior hydration of A β fibrils comes from differential scanning calorimetry (38). However, ssNMR data do not indicate large structural differences between lyophilized and wet fibrils (11,37). Interestingly, experimental studies showed that both A β_{1-40} peptides (52,53) and A β_{1-42} peptides (54) can form ion channels through lipid membranes. However, such A β pores (55) likely occur at a larger scale between *trans*-membrane bundles of protofilaments. Recent simulations of fibrils with up to 32 A β_{16-22} peptides in antiparallel β -strands have also shown similar water channels hydrating the K16/E22 side chains (56).

Stability of connecting loops

A recent study suggested that the loops connecting the N- and C-terminal strands play an important role in aggregation (19,57). To explore the relative contributions of the loop and hydrophobic core to the overall stability and fibril dissociation, we mutated the residues V24–N27 in the salt-bridge region from VGSN to AGAA.

Fig. 5 shows a plot of the C_{α} root mean-square distance (RMSD) of individual residues at 298 K and 398 K. RMSD values were calculated for the interior eight $A\beta_{9-40}$ peptides of finite $S2-C_{22}$ protofilaments, and averaged over the 5–10 ns trajectory segments. At room temperature, we observe only a small increase in the flexibility of the loop region and no significant change in the β -strand segments. At elevated temperature (398 K), the mutations result not only in increased loop flexibility, but also produce large fluctuations in the β -strand regions. Figs. S6 and S7 show snapshots along the trajectories for both wild-type and mutated systems. Overall, the fibril does not appear to be strongly affected by these loop mutations at room temperature, but the enhanced flexibility of the mutated loops results in a significant loss of structure at 398 K.

Conformational dynamics at fibril ends

At elevated temperature, structure is lost most rapidly at the fibril ends, followed by the N-terminal sheets of the interior peptides. The core of the fibrils comprised of the C-terminal β -sheets remains largely intact. The loss of structure is more pronounced in the S2 and S3 systems with larger initial staggering, indicating that (at least at the finite lengths studied here) they are less thermo-stable than S1. The dissociation of peptides from the fibril ends seems to initiate at the N-terminal residues and in the loop region. At 398 K, transient helical conformations form in the loop and the N-terminal regions (emphasized by *boxes* in Fig. 3 B and in Fig. S2 B). Transient helices have also been detected during early fibril formation experiments and simulations (19,22,49,58).

At 398 K, in several instances we observe an exchange in the salt bridge partners from the initial D23/K28 pair to E22/K28 as the end peptides begin to dissociate from the fibril (Fig. 6). The presence of E22/K28 salt bridges has been suggested both for the small peptide aggregates implicated in cytotoxicity and for $A\beta$ monomers in solution (10,19,28,33), but not $A\beta_{1-40}$ fibrils (36).

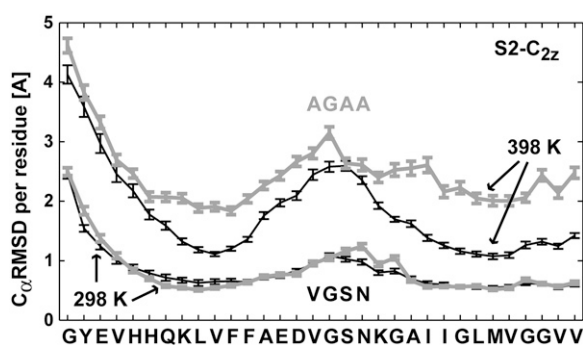


FIGURE 5 C_{α} RMSD per residue calculated for two-peptide units from the interior of $S2-C_{22}$ fibril segments, at 298 K and 398 K, for the wild-type (solid, thin) and a mutated (shaded, thick) loop sequence: $V_{24}GSN_{27} \rightarrow A_{24}GAA_{27}$. Error bars indicate one estimated standard deviation of the average C_{α} -RMSD values.

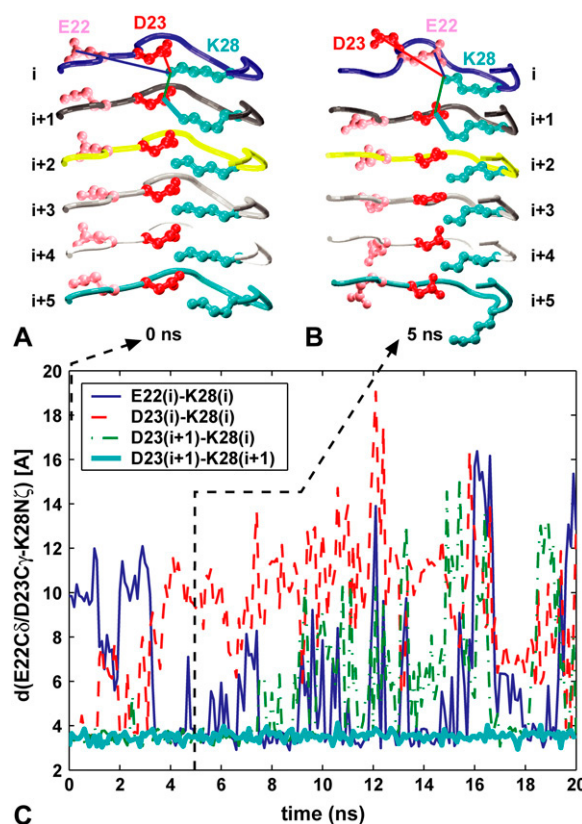


FIGURE 6 Salt bridges (A) D23/K28 and (B) E22/K28 at fibril ends. (C) Time evolution of D23/K28 and E22/K28 distances in the S1 system at 398 K, color-coded as in panels A and B.

Fibril dissociation/elongation

Fig. 7 shows C_{α} RMSD values calculated for a typical trajectory (system S1- C_{22}) at room temperature (298 K) and at 398 K. The RMSD is calculated by using only the C_{α} atoms in the four interior two-peptide units in the $A\beta_{9-40}$ fibril segments, and by aligning the simulation frames to the average obtained for the 1–5 ns trajectory segment. The sequence segments H_5-F_{12} , $E_{14}-G_{21}$, and $I_{23}-G_{30}$ (see Fig. 1 A) were used for the N-terminal sheet, loop, and C-terminal sheet, respectively. Noticeably, the C-terminal β -strands preserve their structural order longer than other regions of the $A\beta$ peptides. C_{α} RMSD values calculated for either the N-terminal sheet or the loop, are always significantly higher, especially at elevated temperatures.

The higher stability of the C-terminal β -strand regions observed in the MD simulations suggests a possible mechanism for fibril elongation that follows the reversed steps of dissociation. In the resulting hypothetical scenario, the initial monomer addition at the end of a growing fibril is driven by strong hydrophobic interactions stabilizing the C-terminal β -strands. In a second stage, the less stable N-terminal β -strands would form. In a final step of this fibril-elongation scenario, the more flexible loop with its relatively hydrophilic

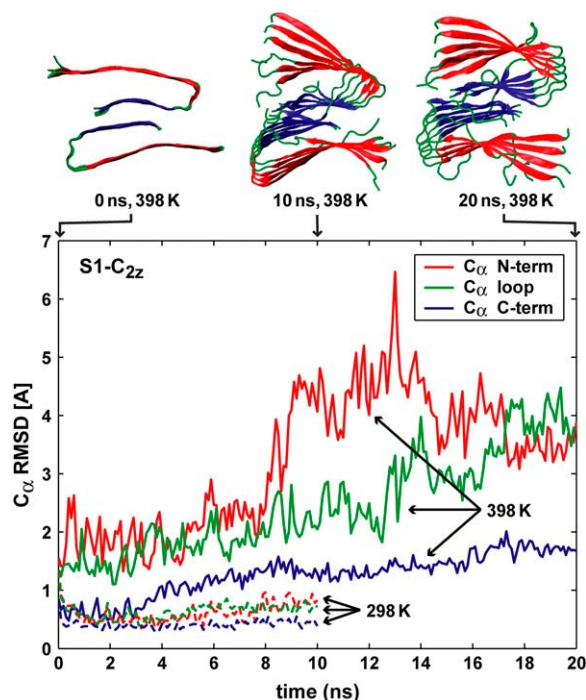


FIGURE 7 C_{α} RMSD values calculated for A β structural elements of the S1-C_{2z} system, as a function of time, with respect to the average conformation of the 1–5 ns trajectory segment at 298 K. The solvent-exposed N-terminal β -strands (red) lose structural order at elevated temperatures faster than loop (green) and C-terminal (blue) regions.

residues would adopt the fibril conformation. During elongation, the E22/K28 ion pair suggested for the free monomer in solution (19,28) is replaced by D23/K28.

Overall, based on our qualitative observations the main driving force for fibril elongation appears to be the formation of C-terminal β -sheets. Their hydrophobic character and matching side-chain motif I31 \times G33 \times M35 \times G37 permits the interdigitation of large and small side chains that stabilizes the quaternary contacts between the two C-terminal β -sheets (16).

Recent experiments showed that perturbing the hydrogen bonds in N- and C-terminal β -sheets through selective N-methylation affects both fibril growth and structure (15). Disrupting the backbone hydrogen bonds of the N-terminal sheets resulted in relatively slow growth of fibrils with a “fuzzy” boundary; with hydrogen bonds in the C-terminal sheets disrupted, the growth was slightly less affected, and the fibrils had a sharply defined surface. These experimental results are consistent with our simulation observations. The C-terminal sheets are held together by strong hydrophobic interactions that would compensate for the partial loss of hydrogen bonds in the N-methylated system and produce wild-type-like fibrils. Increased fibril twisting could be caused by the loss of directional hydrogen bonding, and a gain in importance of packing interactions (59). In contrast, N-methylation at the N-termini disrupts the more fragile

N-terminal sheets, with peptide ends sticking into the solvent to produce fibrils with a “fuzzy” boundary.

CONCLUSIONS

We explored the contributions of the different structural elements of A β protofilaments to the stability, conformational dynamics, and to the fibril elongation/dissociation mechanism. Using multiple 10–20 ns long MD simulations of fibril systems of \sim 70,000 atoms, we studied protofilament models that differ 1), in the relative orientation of the C-terminal β -strands at the fibril core (C_{2x} and C_{2z} , Fig. 1, *D* and *E*); 2), the β -sheet staggering; and 3), connecting loop conformation (S1, S2, and S3 systems, Fig. 2) as well as sequence (VGSN versus AGAA). Our NPT simulations cover a broad range of temperatures (298, 348, and 398 K), for both wild-type and mutated sequences, allowing us to probe the structural stability and the early dissociation events occurring in A β protofilaments. We find that all models are stable at room temperature, and converge toward an interdigitated side-chain packing for intermolecular contacts within and between the two-peptide units of the protofilaments.

The D23/K28 salt bridges maintain a stable and relatively rigid interdigitated structure. However, we find that during the initial stages of fibril dissociation the D23/K28 contacts in A β peptides at fibril ends can break to form the competing E22/K28 interaction. This observation suggests that the loss of E22/K28 contacts could be an important fingerprint of the transition experienced by the A β peptides from their solution structures toward the fibril conformations (19). As reported before (12), we find narrow water channels solvating the D23/K28 salt bridges interior to A β fibril segments.

Simulations at all temperatures reveal that β -strand staggering is a characteristic element of A β protofilaments, permitting a compact, interdigitated packing of side chains from neighboring β -sheets. We find that the type of staggering, as defined by using the intrinsic directionality of A β fibrils, can differ for different fibril models, and we observe a noticeable bias across several fibril models toward adopting positive staggerings (Fig. 2). As the most dramatic change in structure, we observed collective sliding of N- and C-terminal β -sheets on top of each other. Simulations show that A β peptides may adopt structural conformations with smaller β -sheet staggering for residues close to the loop than for those at the tip of the strands (Fig. 4 *D*), with N- and C-terminal strands of the same A β peptide being at an angle to each other. At elevated temperatures, simulations show features common to different models, most notably loss of order in the solvent-exposed N-terminal β -strands coupled to structural disorder in the loop regions. The loop regions and the N-terminal β -strands appear most sensitive to temperature increases. Mutations of loop residues are found to enhance the flexibility of the fibrils at elevated temperatures. Our simulations suggest that the hydrophobic fibril core comprising the C-terminal β -strands of the two molecular

fibril layers is a major stabilizing element and its formation may constitute a crucial step in the aggregation and elongation of A β protofilaments.

SUPPLEMENTARY MATERIAL

An online supplement to this article can be found by visiting BJ Online at <http://www.biophysj.org>.

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