Structural Mapping of Oligomeric Intermediates In An Amyloid Assembly Pathway


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Impact statement:
Solution NMR provides structural and kinetic information about oligomers on pathway to amyloid fibrils that are precisely structured but not cytotoxic.
Abstract

Transient oligomers are commonly formed in the early stages of amyloid assembly. Determining the structure(s) of these species and defining their role(s) in assembly is key to devising new routes to control disease. Here, using a combination of chemical kinetics, NMR spectroscopy and other biophysical methods, we identify and structurally characterize the oligomers required for amyloid assembly of the protein ΔN6, a truncation variant of human β2-microglobulin (β2m) found in amyloid deposits in the joints of patients with dialysis-related amyloidosis. The results reveal an assembly pathway which is initiated by the formation of head-to-head non-toxic dimers and hexamers *en route* to amyloid fibrils. Comparison with inhibitory dimers shows that precise subunit organization determines amyloid assembly, while dynamics in the C-terminal strand hint to the initiation of cross-β structure formation. The results provide a detailed structural view of early amyloid assembly involving structured species that are not cytotoxic.
Introduction

Oligomers have been the focus of amyloid research over decades because of their pivotal role in assembly and their potential cytotoxicity\(^1\). Numerous aggregation-prone proteins (or their fragments) form oligomers\(^2-5\), some of which are cytotoxic\(^6,7\), while others are not\(^8\). Many groups have attempted to elucidate the structure(s) of amyloid oligomers with different biological properties\(^1\). However, their ephemeral nature, dynamic signature, and heterogeneity in mass and conformation provide significant experimental challenges. Hence, our current understanding of the structure of oligomers is often limited to low resolution models\(^7,9-11\), or to oligomers assembled from non-natural amino acids, short peptides, or protein fragments\(^4,5,12\). Establishing a relationship between the oligomers observed and the mechanism of amyloid formation is also an important, but challenging, task. In some cases, oligomers have been shown to be ‘off-pathway’ since they have to dissociate for amyloid formation to proceed\(^13-16\). Characterization of such species, however, does not provide insight into the structural mechanism by which initially unstructured (e.g. A\(\beta\)\(_{42}\), \(\alpha\)-synuclein) or natively structured proteins (e.g. lysozyme, transthyretin, antibody light chains, \(\beta_2\)-microglobulin (\(\beta_2\)m)) undergo conformational conversion to form the parallel in-register cross-\(\beta\) structure of amyloid\(^17\). Other oligomers have been shown to be on-pathway\(^7,11\), or to form via secondary nucleation processes that enhance the rate of fibril formation\(^18\). Proteins in an oligomeric or aggregated form have also been characterized kinetically, thermodynamically and biophysically\(^18-20\). However, a detailed understanding of both the structural properties of oligomers and their role in assembly is needed in order to understand the structural mechanism(s) of amyloid formation and the origins of cytotoxicity, as well as to design inhibitors of the assembly process.

Here we describe an integrative approach which uses kinetic modelling to identify oligomers
formed on-pathway to fibril formation, NMR spectroscopy and other biophysical methods to
determine their structural properties, and cellular assays to determine their cytotoxicity. The
strategy employed can be applied to other assembling protein systems and draws on the
powers of NMR to provide detailed structural information about individual precursors in
dynamic equilibrium within complex mixtures of assembling species, and kinetic modelling
to ascribe their role in amyloid formation. To exemplify this combined structural and kinetic
approach we focus on the naturally occurring variant of human β2m, known as ΔN6. This
variant lacks the N-terminal six amino acids and is formed by natural proteolytic truncation
of the wild-type (WT) protein\textsuperscript{21,22}. WT human β2m (named herein as hβ2m) forms amyloid
deposits in the joints of patients undergoing long term hemodialysis\textsuperscript{23}. However, hβ2m does
not aggregate into amyloid fibrils at physiologically-relevant pH and temperature on an
experimentally accessible timescale \textit{in vitro} (the pH in normal and diseased joints ranges
from 5.5-7.4\textsuperscript{24}). Addition of Cu\textsuperscript{2+} ions, detergents, organic solvents, glycosaminoglycans or
collagen can drive hβ2m amyloid formation at neutral pH\textsuperscript{25-28}. These reagents partially unfold
the native protein, facilitating \textit{cis-trans} isomerization of Pro32 that initiates assembly\textsuperscript{22,25,27,29}.
By contrast with the intransigence of hβ2m to form amyloid \textit{in vitro}, ΔN6 is highly
amyloidogenic, forming fibrils rapidly \textit{in vitro} in the absence of additives at pH 6-7\textsuperscript{21,22,30}.
ΔN6 forms ~30\% of β2m in amyloid plaques in patients with dialysis-related amyloidosis\textsuperscript{31}.
Previous studies have shown that ΔN6 can induce amyloid formation of hβ2m at near-neutral
pH \textit{in vitro}\textsuperscript{22} and can co-assemble with the WT protein into amyloid fibrils\textsuperscript{32}. The structure of
hβ2m in amyloid fibrils formed \textit{in vitro} at low pH (pH 2.0) has also been solved recently
using solid-state NMR and cryo-EM, revealing a parallel in-register cross-β structure typical
of amyloid, which differs dramatically from the all anti-parallel immunoglobulin fold of the
native precursor\textsuperscript{33}. The atomic structure(s) of hβ2m amyloid fibrils formed \textit{in vivo}, and those
of ΔN6 formed \textit{in vitro} or \textit{ex vivo}, however, are not yet known.
Several examples of oligomers (dimers, tetramers and hexamers) of WT hβ2m have been reported previously\textsuperscript{34-42}, with one report of a domain swapped dimer of ΔN6 stabilized by addition of a nanobody\textsuperscript{43}. Since hβ2m is inert to aggregation at physiological pH and temperature \textit{in vitro}, the oligomerization of the protein was stimulated by mutation and/or the addition of Cu\textsuperscript{2+} ions\textsuperscript{34-37}, or by linkage of monomers via non-native disulfide bonds\textsuperscript{38,39}. Although some of these oligomers form under conditions in which WT hβ2m may eventually form fibrils, the role of individual oligomeric species in the aggregation mechanism remains unclear. The oligomers formed in the initiating stages of aggregation of ΔN6 into amyloid also remain obscure.

Here we show that amyloid formation of ΔN6 occurs via a remarkably specific assembly mechanism involving the transient formation of dimers and hexamers. Exploiting NMR methods able to analyze dynamic and lowly-populated states\textsuperscript{44}, we characterize these assemblies, yielding a structural model of the initiating events in ΔN6 aggregation in atomic detail. The results reveal the formation of head-to-head dimers that pack into symmetric hexamers that retain a native-like immunoglobulin fold and are not cytotoxic. The hexamers appear to be primed for further conformational change into the cross-β structure of amyloid by dynamic unfurling of their C-terminal β-strands. The results portray a detailed atomic view of the early stages of ΔN6 assembly that may enable the development of routes to combat disease by targeting the specific protein-protein interactions that define the early stages of assembly.
Results

Fibril elongation occurs through an oligomeric state

Previous results have shown that ΔN6 assembles rapidly into amyloid fibrils in vitro at pH 6.2, but not at pH 8.2\textsuperscript{22}, suggesting that lowering the pH increases the population of aggregation-prone species. Such species may also be relevant in vivo given the acidic microenvironment of the joints of DRA patients\textsuperscript{22,31,42}. At pH 6.2 (close to its pI of 5.8) ΔN6 is dynamic, but retains a native-like immunoglobulin fold\textsuperscript{22}. To determine the kinetic mechanism by which ΔN6 aggregates into amyloid fibrils, experiments were performed in which ΔN6 fibril seeds (20 μM monomer equivalent concentration) were incubated with different concentrations of ΔN6 monomers (20 μM to 500 μM) and the rate of amyloid formation was monitored by the fluorescence of thioflavin T (ThT). All experiments were performed at pH 6.2 at a total ionic strength of 100 mM (see Methods). The simplest kinetic mechanism in which monomers add to the fibril ends would result in a linear dependence of the initial rate of fibril elongation versus the monomer concentration, with saturation at high monomer concentrations\textsuperscript{45-47}. Such behavior is observed for seeded assembly of acid unfolded monomers of hβ\textsubscript{2}m, which initially lack persistent structure\textsuperscript{48}, into amyloid fibrils at pH 2.0 (Figure 1a, b). By contrast, ΔN6 showed more complex behavior, with a clear non-linearity in the initial rate of elongation versus monomer concentration, in which rapid seeded growth occurs only above ~200 μM ΔN6 (Figure 1c, d). This indicates that fibril elongation by ΔN6 must involve addition of one or more oligomeric species to the fibril ends under the conditions employed.

Native-like dimers and hexamers form during ΔN6 assembly

The concentration-dependence of ΔN6 elongation could be explained by an oligomer(s) acting as the elongation unit. To explore whether oligomeric species of ΔN6 are formed
under the conditions employed, sedimentation velocity analytical ultracentrifugation (AUC), size exclusion chromatography (SEC), cross-linking, and NMR experiments were performed. These approaches report on the conformational properties and molecular weight distribution of the assemblies formed at different ΔN6 concentrations. Sedimentation velocity AUC experiments showed that ΔN6 forms discrete oligomers at pH 6.2, with monomers, dimers and higher order species with a sedimentation coefficient (S value) consistent with 6-9-mers (although the rapid equilibration of the species present prevents accurate determination of their mass and population) (Figure 2a). To investigate the molecular mass of the species present, ΔN6 was cross-linked after different incubation times in the absence of fibril seeds using 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride (EDC) (see Methods) and the resulting species examined using SDS-PAGE (Figure 2b). This revealed the presence of hexamers during assembly (Figure 2b). The population of the hexameric species is decreased at later time points, presumably because it is consumed into fibrils (Figure 2b). Analytical SEC of ΔN6 at different protein concentrations without cross-linking revealed only monomers and dimers (Figure 2 - Figure Supplement 1a), consistent with the higher order assemblies dissociating upon dilution on the column. However, when cross-linking was performed prior to SEC, higher molecular weight oligomers were observed, with these species being more abundant when higher protein concentrations were used (Figure 2 - Figure Supplement 1b, c). At the highest concentration of ΔN6 used (500 μM) cross-linking resulted in the rapid formation of high molecular weight aggregates that elute in the void volume (Figure 2 - Figure Supplement 1b, c). The population of these aggregates increases with time, accompanied by depletion of the oligomers, consistent with these species being capable of assembly into amyloid (Figure 2 - Figure Supplement 1d).

$^1$H-NMR and $^1$H-$^{15}$N HSQC NMR spectra of ΔN6 were next acquired to examine the
properties of the oligomers formed. Significant changes in chemical shift and linewidth of individual resonances at different concentrations of ΔN6 were observed, consistent with the finding that ΔN6 self-assembles into higher molecular weight species at pH 6.2 (Figure 2c and Figure 2 - Figure Supplement 2a). The residues most affected lie in the A strand and the BC, DE and FG loops, suggesting that these regions form the intermolecular interfaces in the higher molecular weight species (Figures 2d, e). Consistent with these observations, measurement of the rotational correlation time ($\tau_c$) and diffusion coefficient of the sample, which reflect the average size and shape of the molecules formed, showed a linear dependence on ΔN6 concentration, consistent with protein oligomerization in which the resulting species are in dynamic exchange (Figure 2 - Figure Supplement 1e, f). Together these results show that ΔN6 assembles into dimers and hexamers that are assembly-competent, in dynamic exchange, and assemble via interfaces which are located in the apical region of the protein that surrounds Pro32 (Figure 2e).

To estimate the dissociation constants for dimer and hexamer formation, the chemical shifts and residual dipolar couplings (RDCs) of individual resonances were measured as a function of ΔN6 concentration from 10 - 410 μM (Figure 2 - Figure Supplement 2a-e). Significant chemical shift differences were observed when the ΔN6 concentration was increased from 10 μM to 50 μM without significant line broadening (Figure 2 - Figure Supplement 2c, panels i-iii). Increasing the protein concentration to 100 μM caused a decrease in the chemical shift differences (Figure 2 - Figure Supplement 2c, panel iv), which then increase again in magnitude at 200 μM and 410 μM, accompanied by significant line broadening (Figures 2 - Figure Supplement 2a and 2c, panels v,vi). This complex behaviour is consistent with a monomer-dimer-hexamer equilibrium in which the monomers and dimers have different chemical shifts, while the chemical shifts of dimers and hexamers are similar (an assumption
that is supported by our structural models, see below), and the exchange rate between monomers and hexamers is significantly faster than that between monomers and dimers. Therefore, the monomer-dimer equilibrium dominates the equilibrium (and the observed chemical shift) at low concentrations (50 μM). At higher concentrations the dimer is depleted relative to the hexamer and the chemical shift observed becomes a complex combination of the population of each species, the exchange rate between each species, and the difference in chemical shift of each residue in each assembly. Fitting the chemical shift data to a monomer – dimer – hexamer model yields a $K_d$ for dimer formation of $\leq 50 \mu$M, while that of hexamer formation is $\sim 10 \pm 5 \times 10^{-9}$ M² (see Methods and Figure 2 - Figure Supplement 2b,d), indicating that once dimers form they have a high affinity for one another. Importantly, the monomer – dimer – hexamer model with the estimated affinities adequately describes the observed increase in the $\tau_c$ and the observed diffusion coefficient versus protein concentration (Figure 2 - Figure Supplement 1e, f), independently supporting the model derived. Increasing the $K_d$ for dimer formation to >100 μM results in unrealistically low values for the hexamerization $K_d$ (Figure 2 - Figure Supplement 2d). Moreover, measurement of RDCs versus protein concentration results in a biphasic curve (Figure 2 - Figure Supplement 2e), consistent with a multi-equilibrium process. Using these data the RDCs of the dimer species can be calculated for a range of estimated $K_d$ values (see Methods). Fitting the dimer RDCs to the structure of ΔΝ6 (2XKU), shows significantly poorer fits to the predicted RDC values assuming a dimer $K_d$ higher than 50 μM (Figure 2 - Figure Supplement 2f). To explain the chemical shift and RDC data, therefore, the dimer $K_d$ must be $\leq 50 \mu$M.

**Specific interfaces determine aggregation**

To map the interfaces involved in ΔΝ6 oligomer formation in more detail, intermolecular paramagnetic relaxation enhancement (PRE) experiments were performed. The PRE depends
on the distance between a paramagnet and adjacent nuclei and can provide distance information about (transient) binding interfaces for nuclei that are within ~20 Å of the paramagnetic centre\(^4\), quantified by the effect of the spin label on the relaxation rates of each amide proton (the H\(_N\)-\(\Gamma_2\) PRE rate). \(^{14}\)N-\(\Delta N6\) was spin-labelled with (1-oxyl-2,2,5,5-tetramethyl-D3-pyrroline-3-methyl) methanethiosulfonate (MTSL) by creating Cys variants at positions 20, 33, 54 or 61. Each protein (60 μM) was then mixed with \(^{15}\)N-\(\Delta N6\) (60 μM) at pH 6.2. At this total protein concentration the PREs observed are dominated by the monomer-dimer equilibrium (35% of molecules are monomer, 51% of \(\Delta N6\) molecules are in dimers and 14% of \(\Delta N6\) molecules are in hexamers, determined from the \(K_d\) values measured above). These experiments (Figure 3) showed increased H\(_N\)-\(\Gamma_2\) rates for residues in the A strand and the BC, DE and FG loops when the spin label is attached to residues 33, 54, or 61, suggestive of a head-to-head interaction involving the apical regions of the protein (Figures 3a-c and 3e). In accord with this conclusion, when MTSL is attached at position 20 at the distal side of the protein (Figure 3e), the H\(_N\)-\(\Gamma_2\) rates are vastly decreased (Figure 3d).

To determine whether the head-to-head dimers are critical for aggregation, the AUC, PRE and fibril growth experiments were also performed at pH 8.2 where \(\Delta N6\) does not assemble into amyloid fibrils even after extended incubation times (Figure 3 - Figure Supplement 1a). The sedimentation velocity AUC experiments revealed that monomers and tetramers are formed at pH 8.2, but not hexamers, with the equilibrium in favor of the monomer (Figure 3 - Figure Supplement 1b). Consistent with this, the \(\tau_c\) of 600 μM \(\Delta N6\) at pH 8.2 is ~12 ns, in marked contrast with the \(\tau_c\) of ~50 ns predicted for 600 μM \(\Delta N6\) at pH 6.2 (Figure 2 - Figure Supplement 1e). Finally, intermolecular PRE experiments at pH 8.2 showed small \(\Gamma_2\) rates irrespective of the site of MTSL labelling (Figure 3 - Figure Supplement 1c-d), suggesting that the monomers bind with different affinity and/or via different interfaces at this pH. To
investigate these hypotheses, CPMG relaxation dispersion NMR experiments were performed. These experiments are able to detect excited states populated to as little as 1% of the total protein in solution\textsuperscript{50}. Concentration-dependent CPMG profiles of residues in the B strand, D strand, DE loop, E strand and EF loop were observed at pH 8.2 (Figure 3 - Figure Supplement 2a-d), indicating that the binding interface for tetramer formation differs substantially from the loop-loop interactions in the apical region of the protein that dominate assembly at pH 6.2, despite the fact that ΔN6 retains an immunoglobulin-like fold at both pH values (Figure 3 - Figure Supplements 2e,f). As a consequence of the altered interface that forms at pH 8.2, hexamers and fibrils do not form. Together these results indicate that the head-to-head dimers formed at pH 6.2 are uniquely able to assemble into the hexamers that are crucial for fibril assembly.

**Different dimer structures determine amyloid inhibition and propagation**

To generate dimer structures consistent with the experimental data obtained, simulated annealing molecular dynamics calculations were performed. The calculations converged to two dimer structures (Figure 4a, Figure 4 - Figure Supplement 1 and Table 1). In the lowest energy model (model A) the ΔN6 monomers are arranged in an extended conformation with the N-terminal residues M6 and I7 (WT numbering), along with the BC, DE and FG loops forming the interface (Figure 4a). The inhibitory dimer of ΔN6:murine β\textsubscript{2}m (mβ\textsubscript{2}m) was previously determined using a similar approach\textsuperscript{42}. This dimer also has a head-to-head subunit arrangement, but is characterized by a more acute angle between ΔN6 subunits in which the monomers interact predominantly through the BC and DE loops\textsuperscript{42} (Figure 4b, Video 1). Thus, distinct protein dimers formed from closely related sequences (mβ\textsubscript{2}m and hβ\textsubscript{2}m are 70% identical and 90% similar in sequence) give rise to fundamentally different outcomes of assembly.
Structural models of on-pathway hexamers

Although the majority of the intermolecular PREs can be satisfied by the dimer A structure, the fits are not perfect (Figures 3a-d), presumably since ~14% of ΔN6 molecules form hexamers at the concentration of ΔN6 employed (120 μM). The PRE experiments were thus repeated at higher concentrations (320-400 μM) of ΔN6, wherein > 40% of ΔN6 molecules are predicted to be in hexamers. These experiments revealed a pattern of H_N-Γ_2 rates similar to those obtained at 120 μM ΔN6 (Figure 5 - Figure Supplement 1a-d), with the highest H_N-Γ_2 rates involving the N-terminus, BC, DE and FG loops, suggesting that similar interfaces are formed in the dimer and hexamer. CPMG experiments at 180 μM ΔN6 (26% of ΔN6 molecules are monomer, 48% are dimer, and 26% are hexamer) and 480 μM ΔN6 (13% of ΔN6 molecules are monomer, 32% are dimer and 55% are hexamer) showed that residues in the apical regions of ΔN6, surrounding Pro 32, are also in concentration-dependent exchange at both ΔN6 concentrations at pH 6.2, in support of this conclusion (Figure 5 - Figure Supplement 2).

The ordered nature of assembly (monomer, dimer, hexamer) and the identification of the interfaces involved, allowed us to generate models for the hexameric species (Figure 5 - Figure Supplement 3). The measured PREs were converted into distances and simulated annealing molecular dynamics calculations were performed to create hexamer structures consistent with the experimental PRE and chemical shift data using the lowest energy dimer model (dimer A shown in Figures 4a and 5 - Figure Supplement 3a), as well as the less favourable dimer model B (Figure 4 - Figure Supplement 1e and Figure 5 - Figure Supplement 3b), as starting points. Note that the structure calculation strategy employed does not require knowledge of the dimer and hexamer populations (see Methods). Starting from
dimer A (Figure 4a) the structure calculation resulted in a hexamer in which the three dimers trimerize to form a compact daisy-like structure (Figures 5a-c). The PREs back-calculated from this model are consistent with the experimental data (Figure 5 - Figure Supplement 4). Importantly, hexamer structures generated from dimer B (Figure 4 - Figure Supplement 1e) resulted in poorer fits to the PRE profiles (Methods and Table 2).

In the hexamer models generated from dimer A the dimeric subunits are arranged in a helical manner twisted by ~120°, creating a hexamer that is ~60 Å in diameter and 75 Å in length. This hexamer model is consistent with the collision cross-section (CCS) of ΔN6 hexamers measured using the lowest charge state (15+) (the most native-like species51 (see Methods)) detected using Electrospray Ionization Ion Mobility Spectrometry – Mass Spectrometry (ESI-IMS-MS), but the measured CCS is inconsistent with hexamers derived using dimer B (Figure 5 - Figure Supplement 5a). The monomer-monomer and dimer-dimer interfaces in the best fit hexamer structure (Figure 5a-e) involve similar, but not identical, regions, with the inter-dimer interface extending further into the β-sheet containing the A, B, E and D β-strands, while the intra-dimer interface is formed mostly through the BC and DE loops (Figures 5d-e and Video 2). The formation of dimers generates a hydrophobic surface which becomes buried in the hexamer (Figure 5e, Figure 5 - Figure Supplement 5b and Table 3). Consistent with this, the cross-linked hexamers show a small (1.3-fold) increase in fluorescence in the presence of the hydrophobic dye 8-anilino-1-naphthalenesulfonic acid (ANS), that is much smaller than the ~100-fold increase in ANS fluorescence observed for a typical ‘molten globule’ state52, but similar in magnitude to ANS bound to the highly structured on-pathway folding intermediate of Im7 (monitored using the trapped equilibrium mimic of this species, Im7 L53A54A53 (Figure 5 - Figure Supplement 5d). The ΔN6 dimers show a similar increase in ANS fluorescence as the hexamers despite having a larger exposed
hydrophobic surface area, possibly because ANS binds more weakly or has a lower quantum yield when dimer-bound (Figure 5 - Figure Supplement 5c-d). The interface formed in the inhibitory ΔN6-\(m\beta_2m\) dimer overlaps with the surface required for hexamerization, but not for ΔN6-ΔN6 dimerization (Figure 5f), rationalizing why \(m\beta_2m\) is able to inhibit amyloid formation (note that the \(K_d\) of the \(m\beta_2m:\Delta N6\) complex is 70 \(\mu\text{M}\)\textsuperscript{42}, similar to that (~50 \(\mu\text{M}\)) estimated here for ΔN6 homo-dimerization). The dimers and hexamers were incubated with SH-SY5Y cells, a cell line that is commonly used in studies of amyloid toxicity\textsuperscript{4,7,10,54-56}, and which has been shown previously to take up monomeric and fibrillar \(\beta_2m\)\textsuperscript{54}. Interestingly, there was no evidence for cytotoxicity in assays for 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) reduction, lactate dehydrogenase release, reactive oxygen species formation and cellular ATP level (see Methods) (Figure 5 - Figure Supplement 6). However, rapid dissociation of the uncross-linked oligomers, prevention of conversion to a cytotoxic form by cross-linking, or cytotoxicity requiring different cell types or prolonged exposure (> 24 h) to the oligomers cannot be ruled out.

**Hexamer dynamics may prime further assembly into amyloid**

The hexamer shown in Figure 5 retains a native-like immunoglobulin fold in which the \(\beta\)-strands are anti-parallel. Hence, a major conformational rearrangement has still to occur for ΔN6 to form amyloid fibrils in which the \(\beta\)-strands stack in a parallel in-register structure\textsuperscript{57} (R. Silvers, Y. Su, R.G. Griffin, and S.E. Radford, unpublished). Hints of how this conformational change may be initiated were obtained by quantitative analysis of the CPMG data shown in Figure 6, Figure 5 - Figure Supplement 2 and Figure 6 - Figure Supplement 1. Globally fitting these data for residues which lie in the dimer and/or hexamer interfaces (residues 26, 34, 35, 37, 51, 59, 65, 66, 67, 83, 84, 85, Figures 5d-e) to a two-state fast exchange model yields an exchange rate, \(k_{\text{ex}}^{\text{bind}}\), of 1790 ± 290 s\(^{-1}\) (Figures 6a, b). Distinct
CPMG profiles were observed, however, for residues 87, 89, 91 and 92 which lie in the G strand of monomeric ΔN6 and which are not involved in the dimer-dimer interfaces (they show no significant concentration-dependent chemical shifts, nor are PREs are observed for these residues at low or high protein concentration (Figure 6 - Figure Supplement 1a). The CPMG data for these residues presumably report on conformational changes that result from hexamerization rather than the direct binding event itself. The CPMG data indicate that these residues exchange with a lowly populated (2%) excited state with an interconversion rate, $k_{ex}^G$, of $205 \pm 150$ s$^{-1}$, 10-fold slower than $k_{ex}^{bind}$ (Figures 6c-d and Figure 6 - Figure Supplement 1a-e). Therefore, a distinct process, possibly local unfolding of the C-terminal β-strand, occurs when the hexamer is formed that is driven by the free energy of hexamer formation ($\Delta G_{hexamer}^o \sim 4$ kJ/mol). At ΔN6 concentrations of 480 μM $k_{ex}^G$ is increased to $1170 \pm 196$ s$^{-1}$ (Figure 6 - Figure Supplement 1f-i), consistent with increased hexamer formation enhancing the observed rate of dynamics of the G strand. Hexamer formation thus potentially destabilizes the G-strand of ΔN6, causing local unfolding of this region of the polypeptide chain (although further experiments measuring the sign of the chemical shift change would substantiate this conclusion). This may then lead to more catastrophic structural reorganization of the hexamer into the parallel in-register structure of amyloid (note that the G-strand sequence forms a β-strand in the ΔN6 fibril core$^{58}$). Whether structural conversion occurs within the hexamer, at the fibril end, or requires further, more elaborate molecular steps such as active participation of the fibril surface, or disassembly into smaller structural units prior to fibril assembly, remains to be seen.

**A unified model of ΔN6 polymerization**

As a final test of the validity of the model of ΔN6 assembly proposed we assessed the ability of the structural, kinetic and thermodynamic parameters deduced above to describe the
observed rates of fibril formation measured using ThT fluorescence, as well as the $\tau_c$ values versus $\Delta$N6 concentration measured by NMR, and the fibril yield. Using the dimer and hexamer structural models shown in Figures 4 and 5 and the estimated $K_d$ values for their formation, all of the derived experimental data could be recapitulated (Figure 7). Fitting the seeded fibril growth data to different kinetic models that assume (i) monomers to add to the fibril ends (Figure 7 - Figure Supplement 1a); (ii) monomers are in exchange with a monomeric excited state that is responsible for growth (Figure 7 - Figure Supplement 1b); or (iii) dimers are the elongation units (Figure 7 - Figure Supplement 1c), fail to describe the seeding data (Methods and Table 4). By contrast, a model assuming addition of hexamers describes the ThT kinetic profiles well (Figure 7a), while a more complicated monomer-dimer-tetramer-hexamer model does not improve the fit significantly (Figure 7 - Figure Supplement 1d). The populations of monomer, dimer and hexamer, together with the derived structural models, are also consistent with the observed dependence of $\tau_c$ on protein concentration (Figure 7b). Finally, the amount of hexamer formed (in the absence of seeds) is also predictive of the fibril yield (Figure 7c, d) consistent with the hexamer being required for fibril formation. This conclusion is also supported by the appearance of hexamers early during assembly in the absence of seeds and their disappearance as fibrils form (Figure 2b).
Discussion

Understanding the molecular details of oligomer formation is vital if we are to understand why proteins aggregate into amyloid and why different species have different toxicities\textsuperscript{17,59}. Here we present a general strategy, summarized in Figure 8 - Figure Supplement 1, which allows the identification of oligomeric intermediates in amyloid assembly and enables their structural characterization. By combining the powers of NMR to detect lowly populated species in dynamic exchange, with complementary techniques such as AUC and cross-linking, oligomeric intermediates can be identified and structurally characterized in atomic detail. Importantly, to link these intermediates to the mechanism of aggregation, the derived affinities, stoichiometries and structural models can then be used to globally model the time course of fibril assembly. The strategy presented is not only applicable to protein aggregation, but to any weakly self-associating protein system. Given that the balance between monomers, dimers, higher molecular weight oligomers and fibrils could depend critically on the experimental conditions, including the pH, temperature, protein concentration, amount of seed added, buffer composition and ionic strength, the same protein, or a closely related protein variant, could assemble via different mechanism(s) under different conditions. Indeed, aggregation of many amyloidogenic proteins, including h\beta2m\textsuperscript{33}, is known to result in polymorphic fibrils\textsuperscript{60-63} that could extend via different mechanisms. The approach described here can distinguish between such different assembly pathways and may be able to shed light on the role of individual oligomeric species in aggregation and the origins of amyloid polymorphism.

Using the workflow derived, we show that elongation of ΔN6 amyloid seeds proceeds via a specifically organized hexamer (Figure 8). This finding contrasts with the more common view of monomer addition to fibril ends that has been observed for A\beta_{40/42} (reference\textsuperscript{19}), α-
synuclein\textsuperscript{45}, huntingtin exon 1\textsuperscript{64} and for unfolded h\textbeta_2m at pH 2.0 (Figure 1a)\textsuperscript{65}, while oligomers are thought to play critical roles in the primary/secondary nucleation phases of the assembly of these proteins\textsuperscript{19}. By contrast with these initially disordered proteins, the monomeric precursor of \(\Delta N6\) assembly is structured, a scenario that accounts for more than 20 of the 70 human proteins known to cause amyloid disease\textsuperscript{66}. Other amyloid precursors that are initially structured show an inability to self-seed (e.g. transthyretin\textsuperscript{67}), or display a non-classical dependence of the elongation rate on protein concentration (e.g. light chains\textsuperscript{68}). Whether these and other structured protein precursors assemble by a mechanism akin to that of \(\Delta N6\) could be answered by applying the integrated kinetic and structural approach described here to further examples of this set of proteins.

Here we show that \(\Delta N6\) dimers and hexamers with well-defined interfaces involving the apical regions of the protein are required for fibrils to form under the conditions employed (Figure 8). By contrast, formation of other interfaces, such as that observed here for \(\Delta N6\) at pH 8.2 and the previously reported m\textbeta_2m:\(\Delta N6\) heterodimer\textsuperscript{42} are not able to assemble into amyloid fibrils (Figure 8). The arrangement of subunits in the \(\Delta N6\) dimer and hexamer observed here is different to that in a previously reported structure of a domain swapped \(\Delta N6\) dimer (Figure 8 - Figure Supplement 2c). However, the G strand that is responsible for the domain swap is dynamic in the hexamer structure presented here, consistent with this edge \(\beta\)-strand being able to dissociate from the \(\beta\)-sandwich to form both structures. A variant (H13F) of h\textbeta_2m has also been reported to form hexamers in the presence of Cu\textsuperscript{2+} ions\textsuperscript{34} (Figure 8 - Figure Supplement 2a,b). In the crystal structure of this species the dimers and hexamers interact in a side-to-side or head-to-head manner to create a ring-like assembly, in marked contrast with the daisy-like organization of monomers in the \(\Delta N6\) hexamers shown in Figure 5. Real-time NMR studies of the folding of h\textbeta_2m have also revealed protein concentration-
dependent exchange-broadening in the apical loops of its transient folding intermediate $I_T^{40}$, an observation that has been attributed to head-head oligomers, in agreement with the data presented here for $\Delta N6$ which structurally mimics $I_T^{22}$. The interfaces observed in the $\Delta N6$ dimer and hexamer also differ from the canonical inter-sheet stacking between immunoglobulin domains in antibodies, suggesting that the structural features described here are specific to the dimers and hexamers involved in amyloid assembly. Taken together, the results show that $\beta_2m$ can form different protein-protein interactions, only a specific set of which results in species capable of assembly into amyloid.

Although many studies have attributed the toxicity of amyloid to oligomeric species$^1$, we show here that the dimers and hexamers of $\Delta N6$ are not cytotoxic, at least under the conditions employed, possibly because they are structured and bury substantial hydrophobic surface area. Interestingly, the oligomerization of $\Delta N6$ has been linked to increased toxicity in $C. elegans$ models$^{69}$. Since amyloid formation can proceed via multiple pathways, it is possible that the cytotoxic species of $\Delta N6$ formed in the $C. elegans$ body wall muscle are different to those formed here $in vitro$. For several proteins, cytotoxicity has been ascribed to off-pathway oligomers that accumulate in the lag time of assembly, consistent with amyloid formation being protective for the cell$^{70}$. Interconversion between different forms of oligomers may also be required for cytotoxicity$^7,11$. Such a process could be compromised in the cross-linked species of $\Delta N6$ used here.

In summary, by taking advantage of the power of NMR spectroscopy to visualize transient species, and combining these experiments with detailed analysis of the kinetic, thermodynamic and hydrodynamic properties of the aggregating ensemble of species, we have been able to determine an atomic structural model of two oligomeric species required
for amyloid formation of ΔN6 at pH 6.2, and have generated a model that describes a potential mechanism of fibril elongation from these states. Our findings portray an assembly mechanism that is remarkably well-defined, involving the formation of specific protein-protein interfaces that are unique to the initiating stages of amyloid assembly. Substantial conformational changes have still to occur, however, for the hexameric intermediate to form the cross-β structure of amyloid. How this is achieved remains an open question, but could involve binding to the fibril ends and/or fibril surfaces. Most importantly, the results reveal a remarkable specificity to the early stages of ΔN6 amyloid assembly that involves the formation of well-defined oligomeric species via specific interfaces, the precise details of which determine the course of assembly. These findings suggest new avenues to combat disease by specific targeting of the early intermediates in the amyloid cascade which, at least for ΔN6, involve specific interactions between non-native, assembly-competent states.
Methods

Protein expression and purification

The pINK plasmid containing the ΔN6 gene was transformed into BL21 DE3 plysS E. coli cells. 2 L flasks containing 1 L of LB or HDMI (1 g/L 15N-NH4Cl, 2 g/L 13C-glucose) media were inoculated with 10 mL of starter culture. Cells were incubated at 37 °C, 200 rpm until they reached an OD600 of ~0.6 and then the expression of ΔN6 was induced by the addition of 1 mM isopropyl β-D-1-thiogalactopyranoside (IPTG). Expression was allowed to continue overnight at 37 °C and cells were harvested the next morning using a Heraeus continual action centrifuge at 15000 rpm. The cell pellet containing ΔN6 as inclusion bodies was lysed by the addition of 50-100 mL buffer containing 100 μg/mL lysozyme, 50 μg/mL DNAse I, 50 μg/mL phenylmethanesulfonyl fluoride (PMSF), 10 mM Tris HCl pH 8.2. Further cell disruption was performed using a constant cell disrupter system (Constant Systems) at a pressure of 20.0 kpsi. Inclusion bodies were isolated using centrifugation (15,000 g) for 40 min, 4 °C and the inclusion body pellet was washed with 10 mM Tris HCl pH 8.2 four times. ΔN6 was then solubilized in 10-20 mM Tris HCl pH 8.2 containing 8 M urea (MP Biomedicals) and refolded by dialysis (3000 MW cutoff membrane) against 2-5 L of the same buffer lacking urea. The refolded protein was centrifuged for 30 min at 15,000 g to pellet insoluble material and the supernatant was loaded onto a Q-Sepharose (GE Healthcare) column equilibrated with 20 mM Tris HCl pH 8.2. Bound protein was eluted with a gradient of 0-400 mM NaCl in the same buffer over 800 mL and the protein was freeze-dried after dialysis in 18 MΩ H2O or concentrated using 3000 MW cutoff centrifugal concentrators (Sartorius). The freeze-dried protein was re-suspended in 10 mM sodium phosphate buffer pH 7.0, filtered through 0.2 μm filters (Fisher Scientific) and gel filtered using a HiLoad Superdex-75 Prep column (Amersham Biosciences), calibrated with a standard gel filtration
kit (GE Healthcare). The monomer peak was collected, concentrated, aliquoted and stored at -80 °C or dialyzed into 18 MΩ H₂O and freeze-dried. Cys mutants of ΔN6 were created as described in reference⁴² and purified as above, except that 2 mM dithiothreitol (DTT) was added before gel filtration.

Aggregation assays

ΔN6 seeds were formed by incubation of 800 μM protein in 10 mM sodium phosphate buffer, pH 6.2 containing 83.3 mM NaCl (to give a total ionic strength of 100 mM), 0.02% (w/v) NaN₃ with 200 rpm shaking on a thriller shaker (Peqlab) at 37 °C for 2 weeks. hβ₂m seeds were formed by incubation of 800 μM protein (expressed and purified as described in reference⁴²) in 10 mM sodium phosphate buffer pH 2.0, containing 50 mM NaCl, 0.02% (w/v) NaN₃ with 200 rpm shaking at 37 °C for 2 weeks. The resulting fibrils were sonicated for 15 s to create fibril seeds. For seeding reactions, samples containing 50-500 μM hβ₂m or ΔN6 in pH 2.0 or pH 6.2 buffers, respectively, containing 10 μM thioflavin T (ThT) were incubated quiescently at 37 °C in sealed 96 low binding well plates (Thermo Scientific). De novo ΔN6 fibrils were formed by incubating 60 μM ΔN6 in 10 mM sodium phosphate buffer, pH 6.2, containing 83.3 mM NaCl, 0.02% (w/v) NaN₃ with 600 rpm shaking in a 96 well plate at 37 °C (lag time ~ 20 h) or in an 1.5 mL Eppendorf tube (lag time ~100 h). Control experiments monitoring seeded fibril growth of ΔN6 at pH 8.2 were performed in 10 mM sodium phosphate buffer, pH 8.2 containing 86.6 mM NaCl (total ionic strength 100 mM, identical to that used at pH 6.2) and 0.02% (w/v) NaN₃. Fluorescence was monitored at 480 ± 10 nm after excitation at 440 ± 10 nm using a FLUOROstar Optima micro-plate reader (BMG Labtech).
Analytical ultracentrifugation

For sedimentation velocity experiments, a sample of 450 µL of protein was dialyzed overnight with 10 mM sodium phosphate buffer, pH 6.2 containing 83.3 mM NaCl or 10 mM sodium phosphate buffer, pH 8.2 containing 86.6 mM NaCl (each buffer has a total ionic strength of 100 mM). The sample was inserted in double-sector Epon centerpieces equipped with sapphire windows and inserted in an An60 Ti four-cell rotor. Absorbance data at the appropriate wavelength were acquired at a rotor speed of 48,000 rpm at 25 °C. Data were analyzed using the c(s) continuous distribution of the Lamm equations with the software SEDFIT\textsuperscript{71},

\[ D(s) = \frac{\sqrt{2}}{18} k T s^{\frac{1}{2}} (f/f_0)_w^{3/2} ((1 - \rho_n/\rho) \rho_r^{1/2} ) \]

where \( D(s) \) is the diffusion coefficient, \( k \) Boltzmann’s constant, \( T \) the temperature in K, \( s \) the sedimentation coefficient, \( f \) the frictional coefficient, \( f_0 \) the frictional coefficient of a compact smooth sphere, \( \eta \) the solvent viscosity, \( \rho \) the solvent density and \( \hat{\nu} \) the partial specific volume.

At concentrations over 200 µM 20% of the material sedimented during the initial 3000 rpm run, consistent with the hexamers forming high molecular weight species that sediment before the c(S) data are acquired.

Chemical cross-linking and analytical SEC

ΔN6 (10 µM - 500 µM) was incubated at room temperature without shaking in 10 mM sodium phosphate buffer, pH 6.2 containing 83.3 mM NaCl (total ionic strength of 100 mM), 0.02% (w/v) NaN\(_3\) overnight. A 100-fold molar excess of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride (EDC) (final concentration 1 mM - 50 mM) was added to the reaction, incubated for 10 min with gentle mixing, followed by the addition of 5 mM sulpho-N-hydroxysuccinimide (NHS) for 5 min at room temperature.
Cross-linking was then quenched by the addition of 10-fold molar excess (over the concentration of EDC) of Tris HCl pH 8.0, or for cellular cytotoxicity assays, Dulbecco’s PBS, and samples were then analyzed immediately using an analytical Superdex S75 10/300 GL column (GE Healthcare) equilibrated with the same buffer. A similar protocol was used to cross-link ∆N6 during de novo fibril formation. A 500 μL volume of 80 μM ∆N6 in 10 mM sodium phosphate buffer, pH 6.2 containing 83.3 mM NaCl, 0.02% (w/v) NaN₃ was incubated in a 1.5 mL micro-centrifuge tube at 37°C with 600 rpm vigorous shaking on a thriller shaker. Under these conditions the lag time is ~100 h instead of ~20 h when the protein is incubated in a 96 well plate (Figure 2b and Figure 3 - Figure Supplement 1a). Samples were cross-linked at various time-points during assembly by addition of 8 mM EDC, incubated for 15 min, followed by addition of 5 mM NHS, also incubated at room temperature for 15 min. The cross-linking reaction was quenched by addition of 200 mM ammonium acetate before samples were subjected to analysis by SDS-PAGE. Given the unavoidable dilution of the samples and their re-equilibration during the SEC run, quantitative analysis of the SEC traces of cross-linked and uncross-linked samples was not performed.

Measurement of ∆N6 correlation times

Rotational correlation times (τₐ) of ∆N6 at different concentrations were measured in 10 mM sodium phosphate buffer pH 6.2 containing 83.3 mM NaCl, or the same buffer at pH 8.2 containing 86.6 mM NaCl (total ionic strength for each sample of 100 mM), 25 °C using a ¹H-TRACT experiment with delays of 0.002 – 0.064 s in a Varian Inova NMR spectrometer operating at 750 MHz. At each delay, the signal intensity of resonances in the amide region was integrated and the resulting curve fitted to a single exponential decay function in order to calculate the relaxation rates of the TROSY (Rₐ) and anti-TROSY (Rₛ) spins. The difference
R_\beta - R_\alpha was then converted to the correlation time. Errors were calculated using duplicate measurements.

Diffusion NMR measurements on ΔN6

Diffusion NMR experiments were performed on ΔN6 samples at different concentrations using pulsed field gradient (PFG) NMR spectroscopy using stimulated echoes with bipolar gradients performed on a Bruker Avance III 750 MHz spectrometer. A series of ^1H spectra were collected as a function of gradient strength (g), the signal (S) was integrated and fitted to:

\[ \frac{S}{S_0} = \exp(-d \cdot g^2) \]

where \( S_0 \) is the signal intensity at zero field gradient, \( d \) is the observed decay rate and \( g \) is the strength of the field gradient pulses. The decay rate \( d \) is directly proportional to the diffusion coefficient, \( D \), of the protein.

Chemical shift perturbation and calculation of K_d values

^1H-^15N TROSY spectra of ΔN6 at different concentrations were collected using a 750 MHz Bruker Avance III spectrometer. The combined ^1H and ^15N chemical shift difference was calculated using the function:

\[ \Delta \delta = \sqrt{ (5 \cdot \delta^{1H})^2 + (\delta^{15N})^2 } \]

Chemical shift data at 10 μM, 20 μM, 100 μM, 200 μM and 410 μM ΔN6 were fitted to a monomer (X_1) - dimer (X_2) – hexamer (X_6) model:

\[ X_1 + X_1 \overset{k_i}{\underset{k_i}{\rightleftharpoons}} X_2 + X_2 \overset{k_i}{\underset{k_i}{\rightleftharpoons}} X_6 \]
The equilibrium concentration of hexamer \([X_6]\) was calculated by numerical integration of the above model using scripts written in Python and converted to fractional saturation. The observed chemical shift \((\Delta \delta)\) is then given by:

\[
\Delta \delta = B_{max} \frac{6 \times [X_6]}{[X_1]}
\]

where \(B_{max}\) is the maximum chemical shift difference. To obtain estimates for the monomer-
monomer and dimer-hexamer \(K_d\)s a grid search was performed by fixing the dimer \(K_d\)
\((k'1/k1)\) and the hexamer \(K_d\) \((k'2/k2)\) to different values (Figure 2 - Figure Supplement 2d).

Excellent fits were produced using a dimer \(K_d < \sim50\) µM, while the hexamer \(K_d\) shows a
narrow distribution centered at \(\sim 10 \pm 5 \times 10^{-9}\) M² (Figure 2 - Figure Supplement 2d). To
further validate the estimation of the dimer \(K_d\), RDC experiments were performed as a
function of \(\Delta N_6\) concentration (Figure 2 - Figure Supplement 2e). \(\Delta N_6\) was aligned in 10
mg/mL of PF1 phage (Asla Scientific) and \(H_N\) RDCs were measured using ARTSY. The
biphasic behaviour of the RDCs suggests a three-state equilibrium in agreement with the
monomer-dimer-hexamer model. The first/second transition at lower protein concentration
(blue/pink dashed line in Figure 2 - Figure Supplement 2e) presumably reports on the
monomer-dimer/dimer-hexamer equilibrium, respectively. In order to extract RDCs of the
dimer species the blue dashed line was extrapolated to 100% dimer using various \(K_d\) values.
The resulting data were fitted to the structure of \(\Delta N_6\) in order to calculate the alignment
tensor of the dimer. Using a \(K_d\) greater than 50 µM results in a decrease in the goodness of
the fit (Figure 2 - Figure Supplement 2f), unless a large conformational change in the
monomer is invoked upon dimer formation. However, based on the chemical shift data shown
in Figure 2 - Figure Supplement 2a,c \(\Delta N_6\) remains native-like across all concentrations, thus
placing an upper limit of the dimer \(K_d\) at \(\sim 50\) µM in agreement with the grid search of the
chemical shift data (Figure 2 - Figure Supplement 2d). Note that the calculated tensor
depends highly on the correct RDC values and therefore RDCs were not used in the structure
calculations described below. Chemical shift perturbations for 10 residues that show
significant chemical shift perturbations (11, 12, 23, 26, 50, 51, 52, 67, 68, 97) were fitted
globally to this model, with representative examples shown in Figure 2 - Figure Supplement
2b. Errors on the measured peak positions were calculated as the standard deviation of the
mean for residues that show insignificant chemical shift changes. Errors on the fitted
parameters were computed using Monte Carlo calculations with 100 steps.

To calculate populations of different species, a monomer-dimer-hexamer model was treated
numerically, i.e. the kinetic equations that describe the time-evolution of the concentration of
each species were integrated to $\tau = \infty$, after equilibrium was reached, yielding the equilibrium
concentration (in molar units) of monomers, dimers and hexamers. Since the dimers consist
of two monomers and hexamers of six monomers these concentrations are then converted to
populations (of monomers in the form of dimer or hexamer) using the relationship:

$$p_n = \frac{n[A_n]}{[M_{tot}]}$$

where $n$ is the oligomer order, $[A_n]$ the equilibrium concentration of the oligomeric state and
$[M_{tot}]$ the total protein concentration. The overall rate of assembly, $k_{on}^{over}$, for this 3-state
model is given by:

$$k_{on}^{over} = \frac{k_1^{app} k_2^{app}}{k_1 + k_2^{app}},$$

where

$$k_1^{app} = 2k_1 [M_{eq}]$$
$$k_2^{app} = 3k_2 [D_{eq}]^2$$

And therefore:

$$k_{on}^{over} = \frac{k_1^{app} k_2^{app}}{k_1 + k_2^{app}} = \frac{6k_1 k_2 [M_{eq}] [D_{eq}]^2}{k_1 + 3k_2 [D_{eq}]^2}$$
The overall $k_{on}^{overall}$ rate of assembly and therefore the total population of oligomers scales linearly as a function of the monomer concentration (see inset in Figure 2-Supplement 1f).

**PRE experiments**

The ΔN6 variants ($^{14}$N-labeled) S20C, S33C, L54C and S61C (1-2 mg/mL) were incubated with 5 mM DTT for 20 min, excess DTT was removed using a PD10 gravity column (GE Healthcare) and the protein was then labeled immediately with MTSL by incubation with a 40-fold molar excess (over the total ΔN6 concentration) of the spin label for 4 h in 25 mM sodium phosphate buffer, pH 7.0 containing 1mM EDTA at room temperature. Excess spin label was removed by gel filtration (PD10 column) in the same buffer. Spin-labelled ΔN6 was used directly or stored at -80 °C. In all cases 100% labelling at a single site was confirmed using ESI-MS. For each PRE experiment MTSL-labelled $^{14}$N-ΔN6 (10 μM -80 μM) was mixed with $^{15}$N-labeled ΔN6 (60 μM -240 μM) and the difference of the $^1$H R$_2$ rates between oxidized and reduced (the latter created by addition of 1 mM ascorbic acid) MTSL-labelled $^{14}$N-ΔN6 was measured. Experiments were performed in 10 mM sodium phosphate buffer, pH 6.2 containing 83.3 mM NaCl or 10 mM sodium phosphate buffer, pH 8.2 containing 86.6 mM NaCl (a total ionic strength of 100 mM at each pH value). Data were recorded at 25 °C using a $^1$H-$^1$N correlation based pulse sequence with 5 or 6 time-points (0.0016 - 0.016 s)$^{19}$ and at least 32 scans per incremental delay, utilizing a Bruker Avance III 750MHz spectrometer equipped with a cryogenic probe. R$_2$ rates were extracted by fitting the relaxation data to single exponentials using in-house scripts. The H$_N$-$\Gamma_2$ rate was then calculated as the difference between the R$_2$ rate in the paramagnetic (R$_2$, para) versus diamagnetic (R$_2$, dia) sample:

$$G_2 = R_{2,\text{para}} - R_{2,\text{dia}}$$
Errors were calculated based on the noise of the experiment. The small PRE signal observed when ΔN6 is modified with MTSL at position 20 can be attributed to non-specific binding of the spin label itself to adjacent protein molecules, since addition of free MTSL resulted in a similar PRE profile (not shown). Thus, data arising from spin-labelled ΔN6 at position 20 were not included in quantitative analysis of the PRE experiments.

15N transverse relaxation dispersion CPMG experiments

15N transverse relaxation dispersion CPMG experiments were performed as described in reference75 using samples dissolved in 10 mM sodium phosphate buffer containing 83.3 or 86.6 mM NaCl to maintain a constant ionic strength of 100 mM at pH 6.2 and 8.2, respectively. Spectra were acquired using a Varian Inova 500 MHz spectrometer using a fixed relaxation delay (τ_{cpmg}) of 48 ms or a Bruker Avance III 750 MHz or 950 MHz spectrometer using a delay of 40 ms. Spectra were processed using NMRPipe76 and R_{2,eff} rates were calculated using:

\[ R_{2,eff} = \frac{(I_x/I_0)}{τ_{cpmg}} \]

where I_x is the peak intensity in each experiment and I_0 is the peak intensity in the reference spectrum (with CPMG train applied). CPMG data from two clusters of residues, one reporting on intermolecular interactions (12 residues) and the second reporting on the dynamics of the G strand (4 residues) (see text) were fitted globally to the Bloch-McConnell equations77 describing a two-state exchanging system using the software package ‘relax’78. The fact that dimer and hexamer interfaces partly overlap, complicates the analysis of the CPMG data at pH 6.2. However, at the concentrations used, where either hexamerization is low (180 μM: 26% monomer, 48% ΔN6 molecules as dimer, 26% ΔN6 molecules as hexamer) or dimerization remains constant (480 μM: 13% monomer, 32% ΔN6 molecules as dimer, 55% ΔN6 molecules as hexamer) good quality fits to a simple two-state model were
obtained. The calculated exchange parameters report on both dimer and hexamer formation. Due to this ambiguity, the apparent exchange rates obtained by fitting the CPMG data were not used in the kinetic modelling of the reaction, but used instead to report on the apparent differences in exchange dynamics of different residues as hexamer formation is enhanced.

Calculation of structural models

Structural models of dimers

Simulated annealing calculations were carried out in XPLOR-NIH. To account for the flexibility of the MTSL moiety, the paramagnetic group was represented as a 5-membered ensemble. The computational strategy employed included three PRE potential terms (arising from S61C-ΔN6, L54C-ΔN6 and S33C-ΔN6) and classic geometry restraints to restrict deviation from bond lengths, angles and dihedral angles. Resonances for which an estimation of the R₂ rate in the presence of the oxidized spin label was not possible were incorporated in the protocol as nOe-type restraints with an upper bound of 11.5 Å and a lower bound of 9 Å. Chemical shift perturbations observed upon binding were incorporated as sparse, highly ambiguous intermolecular distance restraints as described in reference. As chemical shifts can be influenced by numerous factors upon protein-protein interaction, the treatment of the derived data undertaken here results in a loose potential term that is unlikely to bias the structure calculation. Finally, the protocol included a weak radius of gyration restraint (R_{gyr}) calculated as $2.2^*_N^{0.38}$, where N is the number of residues in the complex. R_{gyr} is required in order to prevent bias towards more extended structures and tends to underestimate the true value of the radius of gyration. C2 distance symmetry restraints alongside a non-crystallographic symmetry potential term were also implemented to ensure that the two monomers adopt the same conformation in the dimer. The aforementioned potential terms were then used in a rigid-body energy minimization/simulated annealing in torsion angle
space protocol to minimize the difference between the observed and calculated $\Gamma_2$ rates, starting from random orientations. The first step in the structure calculation consisted of 5000 steps of energy minimization against only the sparse chemical shift restraints, followed by simulated annealing dynamics with all the potential terms active, where the temperature is slowly decreased (3000-25 K) over 4 fs. During the hot phase ($T = 3000$ K) the PRE and nOe terms were underweighted to allow the proteins to sample a large conformational space and they were geometrically increased during the cooling phase. Proteins were treated as rigid bodies until the initiation of the cooling phase, where side chains were allowed to float (semi-rigid body calculation). The final step included torsion angle minimization using all potential terms. The calculations converged to two dimer structures shown in Figures 4a (lowest energy, termed dimer A) and Figure 4 - Figure Supplement 1e (dimer B). Both dimers show a head–head configuration with dimer B showing a larger interface which extends from the BC and DE loops to the B and E strands. Fits to the PRE data are of lower quality for dimer B as judged by visual inspection of the fits and the restraints violation (RMS) (Table 1). However, both dimers were used as initial building blocks for calculation of the hexamer models.

Structural models of hexamers

Starting from dimer A or dimer B an initial docking run was performed. Dimers were treated as rigid bodies and placed at random positions. Residues for which chemical shift differences were observed at high protein concentrations were used as sparse distance restraints alongside a geometry energy potential. Three-fold symmetry was imposed together with a non-crystallographic symmetry potential. The energy arising from the four potential energy terms was minimized in order to generate 1000 hexamer structures. The PRE potential energy was not used during the calculation but only in the scoring of the structures generated (together with the energy of the other four terms). Starting from dimer A, the plot of energy
versus RMSD (to the lowest energy structure) (Figure 5 - Figure Supplement 3a) shows the expected funnel shape with 44 of the 50 lowest energy structures sharing a backbone RMSD of 2-3 Å, indicating that these models are close to a structure that satisfies the PRE restraints.

On the other hand, the 50 lowest energy hexamers built form dimer B show an RMSD of up to 35 Å with three clusters formed (Figure 5 - Figure Supplement 3b). Therefore, these four hexamer structures (one arising from dimer A and three from dimer B) were taken forward for the next round of the protocol which consisted of an exhaustive simulated annealing calculation. Since it is difficult to define the extent to which the PREs arise from the dimer and hexamer, the PREs restraints were converted to distance restraints. Residues that show high PREs such that no peak was observed in the spectrum with oxidized MTSL were given no lower bound, while residues not affected by MTSL had no upper bound. This strategy removed some of the dimer – hexamer ambiguity and instead the protocol searched for hexamers that generally interact in the areas which show increased $\Gamma_2$ rates at high $\Delta N6$ concentrations, rather than quantitatively fitting the PRE data. The details of the simulated annealing run were similar to that performed to calculate the dimer structure, but included a three-fold (instead of two-fold) distance symmetry potential term (giving rise to hexamers with a D3 overall symmetry). The final stage of the protocol consisted of refinement in explicit water using XPLOR-NIH. Distances were converted back to PREs to allow comparison with the measured PRE data. Following this protocol, the hexamers produced from dimer A show increased PRE rates in the A strand and BC, DE loops in agreement with the PRE data (Figure 5 - Figure Supplement 4). On the other hand, all hexamers assembled from dimer B show calculated PREs which describe the measured PREs less well (Table 2) (these fits are available on the University of Leeds publicly available library [https://doi.org/10.5518/329](https://doi.org/10.5518/329)). Note that dimer and hexamer models were generated and selected based only on the agreement with the NMR data. Cross-sections of the oligomers
obtained from other experiments were used only as a check of consistency with the models determined. PDBs of the dimers and hexamers have been deposited in the University of Leeds publicly available library (https://doi.org/10.5518/329). The buried surface areas of dimers and hexamers were calculated using the program NACCESS\textsuperscript{82} which calculates the per residue accessible surface area (ASA) given a structural model. A cutoff of 10% loss in ASA between monomers and dimers/hexamers was used.

Native ESI-IMS-MS

ΔN6 samples were exchanged into a buffer consisting of 50 mM ammonium acetate, 50 mM ammonium bicarbonate pH 7.4 using Zeba spin desalting columns (Thermo Scientific) immediately before MS analysis. NanoESI–IMS–MS spectra were acquired using a Synapt HDMS mass spectrometer (Waters) with platinum/gold-plated borosilicate capillaries prepared in house. Typical instrument parameters were: capillary voltage, 1.2-1.6 kV; cone voltage, 40 V; trap collision voltage, 6 V; transfer collision voltage, 10 V; trap DC bias, 20 V; backing pressure, 4.5 mbar; IMS gas pressure, 0.5 mbar; traveling wave height, 7 V; and traveling wave velocity, 250 ms\textsuperscript{-1}. Data were processed with MassLynx v4.1 and Driftscope 2.5 (Waters). Collision cross sections (CCSs) were estimated through a calibration approach using arrival-time data for ions with known CCS values (β-lactoglobulin A, avidin, concanavalin A and yeast alcohol dehydrogenase, all from Sigma Aldrich). Estimated modal CCSs are shown. Theoretical CCSs were calculated for hexameric model structures using the scaled projection approximation method implemented in IMPACT\textsuperscript{83} after performing in vacuo molecular dynamics simulations to account for structural alterations arising from transfer into the gas-phase, as previously described\textsuperscript{84}. Note that the best scoring model agrees with the CCS of the lowest charge state (15+) (which is considered to be most native\textsuperscript{51}) of the hexamer derived independently using the NMR data.
alone. The IMS-MS experiments thus serve as an independent validation of the structural model derived.

**ANS binding**

The ability of different ΔN6 species to bind 8-anilinonaphthalene-1-sulfonic acid (ANS) was measured by mixing 50 µL of each fraction obtained from analytical SEC of 1 µM ΔN6 (see above) with 200 µL of ANS to yield a final concentration of ANS of 200 µM. Fluorescence spectra were recorded using a ClarioStar plate reader (BMG Labtech) using an excitation wavelength of 370 nm and emission from 400 - 600 nm. The concentration of protein used was estimated to be ~240 µM (monomer), 3 µM (dimer) and 1 µM (hexamer). Experiments on Im7 L53A/L54A were performed as described in reference\(^53\).

**Cytotoxicity assays**

ΔN6 (240 µM) was cross-linked with EDC/NHS as described above. 500 µL of cross-linked material was resolved using a Superdex 75 analytical gel filtration column (GE Healthcare) using Dulbecco’s PBS as a mobile phase (Sigma #D8537). 1 mL fractions were collected. SH-SY5Y cells were obtained from an authenticated and mycoplasma free source (ATCC CRL-2266) and were passaged up to 10 times. The cells were mycoplasma tested and found to be negative. The cells were cultured as described previously\(^47\) using 15,000 cells per well in 96-well plates (Corning #3595) for 24 h in 100 µl of growth medium. This time point has been widely used in other studies of cytotoxicity and hence allows comparison of the results obtained with observations on β2m and other amyloid systems\(^7,10,47,55,56\).

Cells were then incubated with 50 µL of each fraction from SEC for 24 h before analyzing cell viability. PBS alone was used as negative control and 0.02% (w/v) NaN₃ was added as a
positive control for cell death. Each experiment consisted of at least three repeats from two
independent cross-linking reactions. The neuroblastoma cell line SH-SY5Y was chosen for
our assays, as this cell line is a widely accepted model for the study of amyloid toxicity and
has been used by other laboratories for β2m and other amyloid forming sequences7,10,47,55,56.

For MTT assays, 10 µL of a 10 mg/mL solution of MTT (Sigma-Aldrich) was added to each
well for 1.5 h. Cell growth media and excess MTT were then removed and reduced MTT was
solubilized using 50 µL DMSO per well. The absorbance of MTT was determined using a
ClarioStar plate reader (BMG Labtech) at 570 nm with background subtraction at 650 nm.
MTT reduction was calculated as a percentage of PBS buffer treated controls (100%) and
cells treated with 0.02% (w/v) NaN₃ (0%).

Cellular ATP was measured using the ATPlite Luminescence ATP detection assay
(#6016963 Perkin Elmer) according to the manufacturer’s protocol. Luminescence was
measured on a PolarStar OPTIMA plate reader (BMG Labtech). Cellular ATP was calculated
as a percentage of PBS buffer treated controls (100%) and cells treated with 0.02% (w/v)
NaN₃ (0%).

Lactate dehydrogenase (LDH) release was measured using a Pierce LDH cytotoxicity assay
kit (#88953 ThermoFisher Scientific) according to the manufacturer’s instructions.
Absorbance was determined using a ClarioStar plate reader (BMG Labtech) at 490 nm with
background subtraction at 680 nm. LDH release was calculated and normalised to detergent
lysed cells (100%) and PBS buffer treated controls (0%).

Reactive oxygen species (ROS) production was determined using 10 µM 2’,7’-
dichlorohydrofluorescein diacetate (H$_2$DCFDA) (#D399 ThermoFisher Scientific). Cells were incubated with H$_2$DCFDA for 10 min prior to the addition of ΔN6 samples from SEC. Fluorescence was recorded after further incubation for 45 min using a ClarioStar plate reader (BMG Labtech) at 540 nm. ROS production was calculated as a percentage of PBS buffer treated controls (100%) and cells treated with 0.02% (w/v) NaN$_3$ (0%). 10 μM H$_2$O$_2$ was used as a positive control for the induction of ROS production and resulted in a 373±21% ROS assay signal compared with incubation with PBS. Each experiment consisted typically of two-to-three independent experiments each containing five replicates per condition. The error bars represent mean S.E, * p 0.05. Raw data are available at [https://doi.org/10.5518/329](https://doi.org/10.5518/329).

**Kinetic modelling of the rates of amyloid formation**

The fibril growth kinetics for ΔN6 in the presence of ΔN6 fibril seeds shown in Figure 1c,d were fitted to five different kinetic models which consisted of two distinct modules (pre-polymerization and polymerization). In model (1) monomers are assumed to add to the fibril ends (this model contains two parameters, the elongation rate, $k_e$, and a fibril depolymerization rate, $k_e'$). In model (2) the monomers are assumed to be in conformational exchange with a monomeric excited state that is responsible for elongation. Model (3) includes a monomer-dimer equilibrium followed by dimer addition to the fibril ends. Models (2) and (3) contain four parameters, monomer-monomer binding/unbinding rates ($k_1$ and $k_1'$) and monomer conformational exchange rates ($k_e$, $k_e'$). In the fourth model (4) a monomer-dimer-tetramer-hexamer equilibrium was considered. Finally, in model (5) a monomer-dimer-hexamer equilibrium was considered. This model contains six parameters, $k_{l1}$, $k_{l1}'$ (monomer-monomer binding), $k_2$, $k_2'$ (dimer-dimer-dimer binding) and $k_e$, $k_e'$ (exchange). The rate equations for all models are listed in Table 4 and were solved numerically using in house scripts written in Python. In the polymerization module, that describes the addition of the
elongation unit (X) to the already formed fibrils, each assembly step was modelled explicitly

(Table 4). The primary output of each model is the mass fraction of each species as a function

of time. To convert the output of the program to ThT fluorescence curves, the mass of the

elongated seeds was multiplied by a fluorescence factor ($K_{th}$). Elongated seeds were assumed
to be any species ($F_i$) that contain more monomers than the preformed seeds added in the

assay ($F_0$) ($1 \leq i \leq N$), where $N$ represents the number of monomers in the fibril at the end of

the reaction. The mass fraction of monomers present in a fibril was assumed to scale linearly

with ThT fluorescence, giving the following expression for calculating the progress curves:

$$F_i(t) = \sum_{i=1}^{N} i [F_i] K_{th}$$

Seeding data using all five starting ΔN6 monomer concentrations were fitted to each model

globally sharing all rate constants using $N=200$ (200 monomers in a fibril which would

correspond to a fibril roughly 500 nm in length[85]). The monomer-dimer $K_d$ value ($k_1' / k_1$) was

fixed to 50 μM. Fitting the kinetic data to the hexamer addition model produces a hexamer $K_d$
of $\sim 1.9 \times 10^{-9}$ M$^2$ similar to the value of $\sim 10 \pm 5 \times 10^{-9}$ M$^2$ $K_d$ obtained by fitting the chemical

shift perturbation data on protein concentration, confirming the robustness of the model and

the approach employed. Using the estimated $K_d$ values to obtain the populations of dimer and

hexamer ($P_{dim}$, $P_{hex}$) and the structural models to calculate correlation times of the dimers and

hexamers ($\tau_{c,dim}$, $\tau_{c,hex}$), the apparent correlation time at each ΔN6 concentration ($\tau_{c,app}$)

(computed as $\tau_{c,app} = P_{mon} \tau_{c,mon} + P_{dim} \tau_{c,dim} + P_{hex} \tau_{c,hex}$, where $P_{mon/dim/hex}$ is the population of
dimer/hexamer, respectively and $\tau_{c,mon/dim/hex}$ is the correlation time of each species (9.8, 18.5

and 60.3 ns, respectively) calculated using the structural models by HYDROPRO[86] matches

the NMR measured $\tau_c$ versus ΔN6 concentration (Figure 7b).

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Data availability

Data are freely available at the University of Leeds depository (https://doi.org/10.5518/329).

Competing Financial interests

The authors have no competing financial interests.
References


compaction of biomolecules upon transition to the gas-phase using ESI-TWIMS-MS. 


Comparison of productive and inhibitory dimers. The ΔN6 subunit in each dimer (this study) is shown as a dark blue cartoon, while the second ΔN6 monomer in the productive dimer and the mβ2m subunit in the inhibitory dimer are shown as light blue and red, respectively. The BC, DE, FG loops are colored in magenta, green and yellow, respectively, while the intra-dimer interface residues are shown as sticks on both subunits.

ΔN6 assembles into dimers and hexamers. The two ΔN6 subunits in the dimer (dimer A) are shown as blue cartoon and grey cartoon/transparent space-filling representations, respectively. The BC, DE and FG loops are colored magenta, green and yellow, respectively. The intra-dimer interface residues are shown as sticks on one subunit and as orange transparent spheres on the second subunit. The hexamer assembly is then shown as a space-filling model, with dimer 1 shown in dark blue/light blue, dimer 2 in dark yellow/light yellow and dimer 3 in magenta/pink. In the last part of the video only dimer 1 is shown as spheres while dimers 2 and 3 are shown as transparent cartoons. The intra-dimer interface is shown in green and the inter-dimer interface is shown in red.
Figure 1: Dependence of the fibril elongation rate on the concentration of soluble protein. Seeded elongation assays for (a) hβ2m at pH 2.0 monitored by ThT fluorescence. 20 μM of preformed seeds of hβ2m (formed at pH 2.0) and varying amounts of soluble protein were added, as indicated in the key. Note that the protein does not aggregate under these conditions in the absence of seeds on this timescale. The dashed line shows the initial rate of each reaction. (b) The initial rate of fibril elongation (shown in units of ThT fluorescence (a.u.)/h) versus the concentration of hβ2m added. The dashed line represents a prediction using a monomer addition model (see Table 4). (c) Seeded elongation assays for ΔN6 using 20 μM preformed seeds formed from ΔN6 at pH 6.2 as a function of the concentration of soluble ΔN6 added. Open blue symbols denote the ThT fluorescence signal of 500 μM ΔN6.
in the absence of seeds. The dashed line shows the initial rate of each reaction. (d) The initial rate of fibril elongation (shown in units of ThT fluorescence (a.u)/h)) versus the concentration of soluble ΔN6 added. The dashed line shows the dependence of the elongation rate (in units of ThT fluorescence (a.u)/h) on the concentration of monomer assuming a monomer addition model (see Table 4). The elongation rate for monomer addition shows a hyperbolic behavior as a function of monomer concentration, with a linear dependence at lower monomer concentrations, followed by a saturation phase at higher monomer concentrations. The simulation in (b) (dashed line) uses a slower microscopic elongation rate ($k_e$) (Table 4) than that used in panel (d) and therefore saturation is not achieved by 410 μM protein in (b), but is in (d). Five replicates are shown for each protein concentration. Error bars show the standard deviation between all replicates.
**Figure 2: ΔN6 oligomer formation.** (a) Sedimentation velocity AUC of ΔN6 at different concentrations, as indicated by the key. Note that the higher order species decrease in intensity at high protein concentrations (>200 μM) consistent with the formation of large aggregates that sediment rapidly before detection (see also Figure 2 - Figure Supplement 1d). (b) SDS PAGE of cross-linked ΔN6 (80 μM) at different time-points during de novo fibril assembly in the absence of fibril seeds (see Methods). Note that dimers are not observed, presumably as they are not resilient to the vigorous agitation conditions used to accelerate fibril formation in these unseeded reactions, or are not efficiently cross-linked by EDC under
the conditions used (see Methods). A negative stain electron micrograph of ΔN6 after 100 h of incubation is shown below. Scale bar – 500 nm. (c) The methyl region of the 1H NMR spectrum of ΔN6 at 400 μM (left) or 10 μM ΔN6 (right). (d) Per residue combined 1H-15N chemical shift differences between the 1H-15N HSQC spectrum of ΔN6 at 10 μM and 400 μM. Blue dots represent residues for which assignments are missing in both spectra. The dashed line represents one standard deviation (σ) of chemical shifts across the entire dataset. Residues that show chemical shift differences >1σ are shown in yellow, >2σ are colored red, and residues for which the chemical shift difference is not significant (<1σ) are colored grey. Residues that are broadened beyond detection in the spectrum obtained at 400 μM are colored in magenta (see also Figure 2 - Figure Supplement 2a). Residues are numbered according to the sequence of the WT protein. Arg 97 is hydrogen bonded to residues in the N-terminus and presumably is indirectly affected by the interaction. (e) The structure of ΔN6 (2XKU) colored in the same scheme as (d). Pro32 is shown in blue space-fill. The buffer used in all experiments was 10 mM sodium phosphate pH 6.2 containing 83.3 mM NaCl (to maintain a constant ionic strength of 100 mM for all experiments), 25 °C.
Figure 2 - Figure Supplement 1: Analysis of ΔΝ6 oligomerization. (a) Analytical SEC traces of uncross-linked ΔΝ6 at different concentrations as indicated in the key. (b) Analytical SEC traces of cross-linked ΔΝ6. (c) Zoom-in of the SEC traces shown in (b). The elution profile of protein standards is shaded in the background. (d) Analytical SEC traces of 500 μM ΔΝ6 0 h (black), 4 h (green) or 24 h (blue) after cross-linking was performed. (e) Protein correlation times ($\tau_c$) measured using a $^1$H-TRACT experiment (see Methods) as a function of ΔΝ6 concentration at pH 6.2, colored as in (b). The black line represents a linear fit to the data. The correlation time of 600 μM ΔΝ6 at pH 8.2 is shown in blue. (f) The exponential decay rate (d) of the $^1$H NMR signal in a diffusion measurement using stimulated echoes as a function of ΔΝ6 concentration. The black line represents a linear fit to the data. The linear scaling of $\tau_c$ and d is predicted from the linear dependence of the overall assembly rate $k^{\text{on}}$ on ΔΝ6 concentration using the calculated $K_d$s and a monomer-dimer-hexamer model (inset) (see Methods). Data points are colored as in (b). Error bars in (e) and (f) are calculated from the noise level of the spectra and are smaller than the marker points.
**Figure 2 - Figure Supplement 2: Estimation of dimer and hexamer Kd values.** (a) $^1$H-$^1$5N HSQC spectra of 10 μM (green) or 400 μM (pink) ΔN6. Resonances which are broadened >80% at 400 μM ΔN6 are indicated on the spectrum. (b) The combined $^1$H-$^1$5N chemical shift differences that report on hexamer formation as a function of ΔN6 concentration (the data at 50 μM are excluded since at this concentration the equilibrium is dominated by dimer formation). The solid lines represent fits to a monomer-dimer-hexamer model using a dimer...
K<sub>d</sub> of 50 μM and a hexamer K<sub>d</sub> of 10 x 10<sup>-9</sup> M<sup>2</sup> (see Methods). Error bars represent the standard deviation of the resonances that do not show significant chemical shift changes between 10 and 410 μM ΔN6. Data were acquired at 25 °C in 10 mM sodium phosphate pH 6.2 containing 83.3 mM NaCl (total ionic strength of 100 mM). (c) <sup>1</sup>H-<sup>15</sup>N HSQC spectra of 10 μM (panel i), 20 μM (panel ii), 50 μM (panel iii), 100 μM (panel iv), 200 μM (panel v) or 410 μM (panel vi) ΔN6. Residues are labelled in panel (i) according to the color scheme of Figure 2d. (d) Reduced χ<sup>2</sup> surface produced by fits to the monomer-dimer-hexamer model using the 10 residues (11, 12, 23, 26, 50, 51, 52, 67, 68, 97) that showed the largest chemical shift changes. (e) H<sub>N</sub> RDCs as a function of the ΔN6 concentration (a single example for A15 is shown for clarity). (f) Reduced χ<sup>2</sup> values for the fitting of RDC data over the 41 residues measured to the structure of ΔN6 as a function of the K<sub>d</sub> value used to extrapolate the RDCs to 100% dimer (see Methods). Error bars were calculated from the noise level of the experiment.
Figure 3: Identification of interacting surfaces in ΔN6 dimers. Intermolecular PRE data for the self-association of ΔN6. $^{15}$N-ΔN6 (60 μM) was mixed with an equal concentration of (a) $^{14}$N-(S33C)ΔN6-MTSL; (b) $^{14}$N-(L54C)ΔN6-MTSL; (c) $^{14}$N-(S61C)ΔN6-MTSL; or (d) $^{14}$N-(S20C)ΔN6-MTSL in 10 mM sodium phosphate buffer, pH 6.2 containing 83.3 mM NaCl (a total ionic strength of 100 mM). The resulting $\Gamma_2$ rates are color-coded according to the amplitude of the PRE effect (see scale bar: grey-insignificant (<20 s$^{-1}$), yellow- >20 s$^{-1}$, red- >50 s$^{-1}$, pH 6.2, 25 °C). Blue dots in the plots are residues for which resonances are not assigned (na) at pH 6.2. Red crosses indicate high $H_N-\Gamma_2$ rates for which an accurate value could not be determined. Control experiments showed that the small PREs arising from $^{14}$N-(S20C)ΔN6-MTSL arise from non-specific interactions with MTSL itself. Solid black lines depict fits to the PRE data for the dimer structure shown in Figure 4a. Note the poor fits for some residues which are sensitive to hexamer formation (14% of ΔN6 molecules) under the conditions used. Residues are numbered according to the WT sequence and the position of β-strands (2XKU22) is marked above each plot. (e) The structure of ΔN6 (2XKU22) with the BC...
loop shown in magenta, the DE loop in green and the FG loop in yellow. The MTSL attachment sites are highlighted as spheres.
Figure 3 - Figure Supplement 1: Lack of a hexamer population precludes aggregation of ΔN6 at pH 8.2. (a) Aggregation assays for 60 μM ΔN6 monitored by ThT fluorescence at pH 6.2 (red) or pH 8.2 (blue), 37 °C with agitation (600 rpm). Five replicates are shown. Negative stain transmission electron micrographs of samples at 100 h are shown alongside in the same color code. (b) Sedimentation velocity AUC traces for 120 μM ΔN6 at pH 6.2 (red) or pH 8.2 (blue). (c,d) Intermolecular PRE values for ΔN6 at pH 8.2. 60 μM 15N-ΔN6 was mixed with (c) 60 μM of 14N-(S61C)ΔN6-MTSL or (d) 60 μM of 14N-(S20C)ΔN6-MTSL in 10 mM sodium phosphate buffer, pH 8.2 containing 86.6mM NaCl (total ionic strength 100 mM). Γ2 rates are color-coded according to their amplitude (blue-not assigned, grey-insignificant (<20 s⁻¹), yellow- >20 s⁻¹ red- >50 s⁻¹ at pH 8.2, 25 °C). Residues are numbered according to the WT sequence. The position of β-strands (from 2XKU) is marked above each plot.
**Figure 3 - Figure Supplement 2: Mapping the interface of ΔN6 self-association at pH 8.2 using CPMG experiments.** $^{15}$N Relaxation dispersion CPMG data for residues (a) V49, (b) E74 and (c) Y78 at 1200 μM (blue) or 600 μM ΔN6 (red). Solid lines represent fits to a fast exchange model (see Methods). (d) Plots of $R_{ex}$ (defined as $R_{2,eff}^{50Hz} - R_{2,eff}^{680Hz}$) per residue at different concentrations of ΔN6 at pH 8.2 as indicated in the key. The dashed line represents one standard deviation of the mean. (e) Correlation plot of $H_N$ RDCs measured at pH 6.2 versus those back-calculated from the structure of ΔN6 (2XKU) at pH 7.5. (f) The structure of ΔN6 monomers (2XKU) colored according the amplitude of the $R_{ex}$ values at 1200 μM shown in (d). The results show that the interface between interacting monomers at pH 8.2 involves interaction between β-sheets mediated by residues in the B, D and E β-
strands and adjacent residues in the DE loop. This interface is very different to the loop-loop interactions that create the dimer interface at pH 6.2 (see Figure 3e).
Figure 4: Structural models of ΔN6 dimers. Structural models of (a) the lowest energy ΔN6 homodimer (dimer A) and (b) the ΔN6-mβ2m heterodimer that inhibits ΔN6 fibril assembly\textsuperscript{42}. Interface residues (identified as those residues that have any pair of atoms closer than 5 Å) are shown in a ball and stick representation on one subunit and are colored in space fill in gold in (a) or red in (b) on the surface of the second subunit. ΔN6 is shown in the same pose (blue) in (a) and (b). The BC, DE and FG loops are shown in magenta, green and yellow, respectively, and the position of attachment of MTS{	extlt;i}L for the PRE experiments (residues 20, 33, 54 and 61) is highlighted in spheres. PDB files are publicly available from the University of Leeds depository (https://doi.org/10.5518/329). See also Video 1.
Figure 4 - Figure Supplement 1: Alternative ΔN6 dimer structures. Intermolecular PRE data for the self-association of $^{15}$N-ΔN6 (60 μM) mixed with 60 μM of (a) $^{14}$N-(S33C)ΔN6-MTSL, (b) $^{14}$N-(L54C)ΔN6-MTSL, (c) $^{14}$N-(S61C)ΔN6-MTSL, or (d) $^{14}$N-(S20C)ΔN6-MTSL) with the PRE effect color-coded according to its amplitude (blue dot- residues not assigned, grey-insignificant (<20 s$^{-1}$), yellow- >20 s$^{-1}$, red- >50 s$^{-1}$, pH 6.2, 25°C). Red crosses indicate high H$_N$-$\Gamma_2$ rates for which an accurate value could not be determined. Solid black lines represent back-calculated PREs from the high energy dimer structure shown in (e).
and (f). The small PREs arising from $^{14}$N-(S20C)ΔN6-MTSL result from non-specific interactions with MTSL itself. (e and f) The structural model of dimer B shown in different orientations. In each diagram one subunit is shown in cartoon representation (BC loop (magenta), DE loop (green) and FG loop (yellow)) and the second is shown as a surface. Interface residues are highlighted as balls and sticks on the first subunit and shown in gold space-fill on the second subunit. The MTSL attachment sites are highlighted as spheres and the positions of attachment (20, 33, 54 or 61) are labeled. The interface in dimer B involves a more extensive inter-subunit interface in the apical loops than observed in dimer A (Figure 4a). The resulting interface for dimer B does not describe the PRE data (solid black line) as well as the lower energy model of dimer A shown in Figure 4a (Methods and Table 1).
Figure 5: Structural model of ΔN6 hexamers. (a-c) Sphere representations of the hexamer model formed from dimer A rotated by 90° in each view. Subunits belonging to the same dimer are colored in different tones of the same color. (d) The monomer-monomer (intra-dimer) interface is highlighted in green on the surface of the dimer formed from subunits 1a and 1b (within dimer A), with the other dimers shown as cartoons. (e) The inter-dimer interface is colored red on the surface of the dimer formed from subunits 1a and 1b, with the
dimers shown as cartoons. (f) As in (e), but showing the dimer formed from subunits la and 1b, superposed with the mβ2m subunit in the inhibitory ΔN6-mβ2m dimer (green cartoon).

The ΔN6-ΔN6 and ΔN6-mβ2m dimers were aligned on the ΔN6 subunit 1b. Schematics of the assemblies are shown at the bottom colored as in (d-f). Note that the BC, DE and FG loops are highlighted as thicker chains in blue, green and cyan, respectively, in d-f. PDB files are publicly available from the University of Leeds depository (https://doi.org/10.5518/329).

See also Video 2.
Figure 5 - Figure Supplement 1: Intermolecular PREs at high ΔN6 concentration.

Intermolecular PRE data for the self-association of (a) 240 μM $^{15}$N-ΔN6 mixed with 80 μM $^{15}$N-(L54C)ΔN6-MTSL, (b) 200 μM $^{15}$N-ΔN6 mixed with 200 μM $^{14}$N-(L54C)ΔN6-MTSL,

or (c) 80 μM $^{15}$N-ΔN6 mixed with 240 μM $^{14}$N-(S33C)ΔN6-MTSL. PRE data are color-coded according to their amplitude (blue dots-not assigned, grey-insignificant (<20 s$^{-1}$), yellow- > 20 s$^{-1}$, red- > 50 s$^{-1}$, pH 6.2, 25 °C). Red crosses indicate high $H_N-\Gamma_2$ rates for which an accurate value could not be determined. (d) Raw PRE data for residue 85V when 60 μM $^{14}$N-(L54C)ΔN6-MTSL was mixed with 60 μM $^{15}$N-ΔN6 (left) or when 200 μM $^{14}$N-(L54C)ΔN6-MTSL was mixed with 200 μM $^{15}$N-ΔN6 (right). Solid lines represent single exponential fits for the paramagnetic (black) or the diamagnetic samples (red).
Figure 5 - Figure Supplement 2: Additional interfaces do not form in the ΔN6 hexamer. 15N relaxation dispersion CPMG data for residues (a) 37, (b) 67, and (c) 83 at 180 μM ΔN6 (26% ΔN6 molecules are monomers, 48% are in dimers, 26% are in hexamers) (red) or 480 μM ΔN6 (13% ΔN6 molecules are monomers, 32% are in dimers, 55% are in hexamers) (black). Solid lines represent fits to the fast exchange model, yielding values of \( k_{\text{ex}}^{\text{bind}} \) of 1790 ± 290 s\(^{-1}\) at 180 μM ΔN6 and \( k_{\text{ex}}^{\text{bind}} \) of 1170 ± 196 s\(^{-1}\) at 480 μM ΔN6 (see Methods). (d) Plots of \( R_{\text{ex}} \) per residue defined as \( R_{\text{2,eff}}^{500\text{Hz}} - R_{\text{2,eff}}^{680\text{Hz}} \). The dashed line represents one standard deviation of the mean calculated for all data points. Residues are numbered according to the WT sequence. Significant CPMG profiles are observed for residues in the N-terminus, A strand, BC, DE and FG loops, in excellent agreement with the intermolecular PRE data shown at 120 μM and 320 μM ΔN6 in Figure 3 and Figure 5 - Figure Supplement 1. Residues which are severely broadened at 480 μM, thereby precluding accurate determination of their \( R_{\text{ex}} \) values, are shown as black crosses. Crucially, when the protein concentration was increased the residues which show significant CPMG profiles are unchanged suggesting that the dimers and hexamers share a similar interface. (e) The
structure of ΔN6 (2XKU^{22}) colored according to the $R_{ex}$ amplitude as indicated in the scale bar. *Trans* Pro32 is shown in space-fill (pale blue).
**Figure 5 - Figure Supplement 3: Initial docking of dimer structures to create hexamer models.** Plots of RMSD (to the lowest energy structure) versus total energy for hexamers generated by docking of (a) the lowest energy dimer structure (dimer A) or (b) the higher energy dimer (dimer B). The 50 lowest energy hexamer structures are marked as red circles. The hexamers that were selected for the next round of structure calculation for each dimer starting model are marked with green arrows. The structural model of dimer A and dimer B are shown alongside colored as in Figure 4 - Figure Supplement 1.
Figure 5 - Figure Supplement 4: Intermolecular PREs back-calculated from the hexamer structural model generated from dimer A. Intermolecular PRE data for the self-association of ΔN6. 15N-ΔN6 (60 μM) was mixed with 60 μM of (a) 14N-(S33C)ΔN6-MTSL, (b) 14N-(L54C)ΔN6-MTSL, (c) 14N-(S61C)ΔN6-MTSL, or (d) 14N-(S20C)ΔN6-MTSL. The data are color-coded according to their amplitude (blue dots-not assigned, grey-insignificant (<20 s⁻¹), yellow- > 20 s⁻¹, red- > 50 s⁻¹, pH 6.2, 25 °C). Red crosses indicate high H_N-Γ_2 rates for which an accurate value could not be determined. Solid black lines represent back-calculated PREs from the lowest energy hexamer structure (arising from dimer A) shown in Figure 5. The RMS distances (Å) between the intermolecular distances that were used as restraints and those back-calculated from the hexamer structural model are shown (inset) for each dataset (see Methods).
Figure 5 - Figure Supplement 5: Conformational and biochemical properties of ΔN6 hexamers. (a) ESI-IMS-MS analysis. Collision cross section (CCS) distributions for each observed charge state of hexameric ΔN6. The charge state for each CCS distribution is indicated. Note that the CCS of the lowest (most native) charge state (15+) is consistent with the hexamer model generated from dimer A (labelled A (green)), but not the models generated from dimer B (labelled (B(i)), (B(ii)) and (B(iii)) for the three conformers labelled in Figure 5 - Figure Supplement 3b). (b) Hydrophobicity of the hexamer interface. The surface of dimer 1 in the hexamer is colored according to the Eisenberg hydrophobicity scale (Arg=-2.53, Ile=1.38) with the other dimers shown as cartoons. A key is show alongside. The view on the left-hand side shows the surface that is packed against dimers 2 and 3 in the hexamer (interior), with the view on the right-hand side showing the exterior surface of the assembly. (c, d) Fluorescence emission spectra of ANS (200 μM) incubated with (c) ΔN6
monomers (green), (d) dimers (open symbols) or hexamers (red) (eluting at 17 mL, 15 mL and 11 mL, respectively obtained with/without cross-linking, as indicated, using SEC (Figure 5 - Figure Supplement 6)). The fluorescence emission spectrum of ANS in buffer alone is shown in blue. ANS bound to the partially folded Im7 variant L53A I54A\textsuperscript{53} (1 μM) is shown for comparison (black). This was used as a model for a compact native-like folding intermediate\textsuperscript{53} (see text).
Figure 5 - Figure Supplement 6: ΔN6 oligomers are not cytotoxic to SH-SY5Y cells.

Toxicity of cross-linked (solid line/grey bars) or uncross-linked (dotted line/white bars) ΔN6 species following purification by analytical SEC. Cell toxicity was assessed using MTT reduction, cellular ATP level, generation of reactive oxygen species (ROS), and LDH release assays. For assays of MTT reduction, ATP levels and ROS production, the data are normalized to PBS (100%) and NaN₃ treated controls (0 %). LDH release is normalized to detergent lysed cells (100 %) and PBS buffer treated controls (0 %). The error bars represent...
mean S.E, * p 0.05. No evidence for cytotoxicity was observed for any protein species under the conditions employed.
Figure 6: G-strand unfurling may occur upon hexamer formation. $^{15}$N CPMG relaxation dispersion data at 750 MHz (magenta) and 950 MHz (red) (180 μM ΔN6, pH 6.2 (26% ΔN6 molecules are monomers, 48% are in dimers, 26% are in hexamers) for residues (a) 51, (b) 37, (c) 89, and (d) 92. Residues 37 and 51 report on intermolecular interactions that describe dimer and/or hexamer formation (schematic, top left), while residues 89 and 92 do not lie in an interface and report instead in the dynamics of the G strand in the different assemblies formed. The position of all five residues used in the cluster analysis of G strand dynamics is shown in spheres on the structure of ΔN6 (blue cartoon, top right). Pro32 is shown as a magenta sphere. Solid lines represent global fits to the Bloch-McConnell equations.
(Methods) for each cluster of residues. The extracted parameters of the global fit for the two processes ($k_{ex}^{\text{bind}}$ and $k_{ex}^{G}$) are indicated above the plots.
Figure 6 - Figure Supplement 1: Hexamer formation increases the dynamics of the G strand. (a) Location of the G strand in relation to the dimer and hexamer interfaces. Dimer 1 in the hexamer is shown in a cartoon representation while dimers 2 and 3 are shown as semi-transparent surfaces. The positions of the amide protons for residues 87, 89, 91, 92 are shown as grey spheres and the residues that take part in both the dimer and hexamer interfaces are shown as red spheres on the structure of dimer 1. A schematic of the assembly is shown alongside. $^{15}$N relaxation dispersion CPMG data for residues (b) 87, (c) 89, (d) 91 and (e) 92 at 950 MHz (red) and 750 MHz (magenta) of 180 $\mu$M $\Delta$N6, pH 6.2. Solid lines represent the global fits to all residues in the cluster to the slow exchange model which yields a $k_{ex}^{G}$ of 205 ± 150 s$^{-1}$. CPMG data for the same residues (f) 87, (g) 89, (h) 91 and (i) 92 at 750 MHz (blue) and 600 MHz (grey) using 480 $\mu$M $\Delta$N6. Solid lines represent global fits to the fast exchange model which yields a $k_{ex}^{G}$ of 1170 ± 196 s$^{-1}$. 
Figure 7: The monomer-dimer-hexamer model describes the thermodynamics and kinetics of fibril elongation. (a) Global fits (blue solid lines) to the fibril elongation kinetics monitored by ThT fluorescence assuming a hexamer addition model at different concentrations of soluble ΔN6 (dots) (Methods and Table 4). The concentrations of ΔN6 are colored according to the key. The average of five replicates is shown. (b) Protein correlation times ($\tau_c$) measured using NMR (red) and back-calculated values (green) using the populations of monomers, dimers and hexamers predicted from the monomer-dimer-hexamer model and the correlation times of the dimers and hexamer structural models shown in Figures 4 and 5. (c) The fibril yield (after 100 h) of each elongation reaction. SDS-PAGE analysis of the whole reaction (shown in (a)) before centrifugation (W) or of the supernatant
(S/N) after centrifugation at the different concentrations of ΔN6, as indicated. (d) Bar-charts showing the % of insoluble material (grey) measured using densitometry of the gel shown in (c). The % hexamer population in the absence of seeds (black) predicted by the monomer-dimer-hexamer model at each ΔN6 concentration correlates with the % insoluble material (grey). Note that the fibril yield is low since fibrils cannot form when the monomer concentration falls significantly below the K_d for dimer formation (50 μM).
Figure 7 - Figure Supplement 1: Alternative kinetic models do not describe the kinetics of seeded fibril growth. Global fits (blue solid lines) which assume that (a) a monomer, (b) a monomer excited state, or (c) a dimer, add to the fibril ends do not describe the observed fibril growth kinetics monitored using ThT fluorescence at different concentrations of soluble ΔN6 (dotted lines and key). A more complex monomer-dimer-tetramer-hexamer model (d) does not improve the quality of the fit compared with that shown in Figure 7a.
Figure 8: Fibril formation in atomic detail. Schematic representation of the mechanism of amyloid formation for ΔN6. During folding of hβ2m, a highly dynamic intermediate with a flexible A strand is populated prior to formation of the native-like intermediate termed ΙΤ, which has a native-like fold but contains a non-native trans X-Pro32 bond. The latter species is mimicked by ΔN6 and formed in vivo by proteolytic degradation of the WT protein. Only ΙΤ/ΔN6 is primed for aggregation, while the intermediate with the flexible A strand is not able to assemble directly into amyloid. As reported here, ΔN6 forms elongated head-to-head dimers (upper image, centre) which assemble into hexamers. Alternative dimers involving interactions between the ABED β-sheets in adjacent molecules formed at pH 8.2 (lower image, centre) do not associate further into fibrils. Murine β2m (mβ2m) also interacts with ΔN6 at pH 6.2 to form head-to-head heterodimers. The subunit orientation is different in this heterodimer, occluding the hexamer interface and inhibiting assembly (central image). ΔN6 hexamers can elongate fibrillar seeds and show enhanced dynamics in the G strand which...
could represent the first step towards the major structural reorganization required to form the parallel in-register amyloid fold. How this final step occurs, however, remains to be solved.
Figure 8 - Figure Supplement 1: A workflow to enable weakly self-assembling systems to be analyzed in structural, kinetic and thermodynamic detail. A schematic overview of the strategy employed to study the aggregation of ΔN6 which can be extended to other systems. Careful examination of kinetic rates of aggregation leads to the identification of possible aggregation pathways. Structural methods (AUC, SEC, cross-linking, ESI-IMS-MS) can then be used to identify the molecular weight and collision cross section of the species involved. NMR chemical shift analysis and measurements of RDCs can be used to determine estimates of $K_d$ which in turn can be used to determine conditions under which different species are populated. More detailed NMR studies lead to structural models of these species, while stabilization of the intermediates by chemical cross-linking aids the assessment of their
cytotoxicity. The structural and kinetic information collected leads to the generation of kinetic models whose ability to describe the progress of aggregation monitored by ThT fluorescence is tested using numerical methods. Agreement is suggestive of the validity of the kinetic mechanism of assembly and the identity and structural properties of oligomeric intermediates formed.
Figure 8 - Figure Supplement 2: Examples of some previously characterized oligomers of WT hβ2m and ΔN6. (a, b) Cu^{2+}-stabilized H13F hβ2m hexamer^{34,88-90}. (c) Domain swapped ΔN6 dimer^{43}. 
Table 1. Agreement between experimental and back-calculated intermolecular PREs for the two dimer structures (dimer A and dimer B (see Figure 5 - Figure Supplement 3). RMS values are shown comparing the measured versus the predicted values from the structure PREs measured from S33, L54 and S61. Data from position S20 were not used as they arise from non-specific interactions with MTSL. *High PREs refer to PREs in the BC loop (measured from S33, L54 and S61) that (due to their large value) could not be measured accurately and therefore are incorporated as loose distance restraints.

<table>
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<th>PRE term</th>
<th>RMS Dimer A</th>
<th>RMS Dimer B</th>
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<tbody>
<tr>
<td>S33C(ΔN6)-ΔN6 (s⁻¹)</td>
<td>18.65</td>
<td>15.10</td>
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<td>L54C(ΔN6)-ΔN6 (s⁻¹)</td>
<td>29.02</td>
<td>27.44</td>
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<tr>
<td>S61C(ΔN6)-ΔN6 (s⁻¹)</td>
<td>19.44</td>
<td>23.27</td>
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<tr>
<td>*High PREs (Å)</td>
<td>2.78</td>
<td>3.79</td>
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Table 2. Agreement between experimental and back-calculated intermolecular distances for different hexamer structures. RMS values are shown comparing the measured versus the predicted distances from each structural model for distances measured from S33, L54 and S61. Data from position S20 were not used as they arise from non-specific interactions with MTSL. See also Figure 5 - Figure Supplement 3.

<table>
<thead>
<tr>
<th>PRE term</th>
<th>Hexamer 1 RMS (Å)</th>
<th>Hexamer 2(i) RMS (Å)</th>
<th>Hexamer 2(ii) RMS (Å)</th>
<th>Hexamer 2(iii) RMS (Å)</th>
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<tbody>
<tr>
<td>S33C(ΔN6)-ΔN6</td>
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<td>2.68</td>
<td>2.58</td>
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<td>L54C(ΔN6)-ΔN6</td>
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<td>2.33</td>
<td>2.26</td>
<td>1.87</td>
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<td>S61C(ΔN6)-ΔN6</td>
<td>2.22</td>
<td>2.7</td>
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<td>3.11</td>
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Table 3: Analysis of dimer and hexamer interfaces. The buried surface area is calculated as the sum for the two subunits for each complex. Interface residues were identified as those residues that lose at least 10% of accessible surface area upon oligomer formation.

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<tr>
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<th>ΔN6 Dimer A</th>
<th>ΔN6 Hexamer</th>
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</thead>
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<tr>
<td>Buried Surface Area (Å²)</td>
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<td>% Charged residues in the interface</td>
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<td>% Hydrophobic residues in the interface</td>
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<td>54</td>
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<tr>
<td>Module</td>
<td>Variant</td>
<td>Reaction scheme</td>
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<td>-------------------------------</td>
<td>--------------------------------</td>
<td>-----------------</td>
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<tr>
<td>Pre-polymerization</td>
<td>No Pre-polymerization (Monomer addition)</td>
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<td></td>
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<tr>
<td></td>
<td>Monomer conformational exchange</td>
<td>$X_1 \overset{k}{\underset{k'_1}{\rightleftharpoons}} X'_1$</td>
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<tr>
<td></td>
<td></td>
<td>$X = X'_1$</td>
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<td>Dimer addition</td>
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<td></td>
<td></td>
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<td></td>
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<td>$X = X_6$</td>
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<tr>
<td></td>
<td>Monomer-Dimer-Tetramer-Hexamer</td>
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Table 4. Reaction schemes, rate equations and rate constants for the fibril elongation models tested. X represents the species that add onto the fibril ends.