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NMR structure of the water soluble $A\beta_{17-34}$ peptide

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Synopsis

Alzheimer's disease is the most common neurodegenerative disorder in the world. Its most significant symptoms are memory loss and decrease in cognition. Alzheimer's disease is characterized by aggregation of two proteins in the brain namely $A\beta$ (amyloid β) and tau. Recent evidence suggests that the interaction of soluble $A\beta$ with nAChR (nicotinic acetylcholine receptors) contributes to disease progression. In this study, we determine the NMR structure of an $A\beta_{17-34}$ peptide solubilized by the addition of two glutamic acids at each terminus. Our results indicate that the $A\beta$ peptide adopts an α -helical structure for residues 19–26 and 28–33. The α -helical structure is broken around residues S26, N27 and K28, which form a kink in the helical conformation. This α -helix was not described earlier in an aqueous solution without organic solvents, and at physiological conditions (pH 7). These data are in agreement with $A\beta$ adopting an α -helical conformation in the membrane before polymerizing into amyloid β -sheets and provide insight into the intermediate state of $A\beta$ in Alzheimer's disease.

Key words: Alzheimer's disease, amyloid beta, amyloid β , $A\beta_{17-34}$, NMR, soluble peptide

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INTRODUCTION

Alzheimer's disease is widely characterized by amyloid peptide deposits inside brain tissues [1]. These deposits, also called amyloid plaques, present one of the most characteristic feature of the disease and can be seen in different brain regions [2]. Amyloid plaques consist of $A\beta$ (amyloid β) peptides that become aggregated [3–5]. The secondary structure of aggregated $A\beta$ is mostly β -sheet [6–8] and results from enzymatic cleavage of the APP (amyloid precursor protein) [9,10] by α , β and γ -secretases [11]. When APP is cleaved by α -secretase, the resulting product is a protein named soluble APP α , while cleavage with the two other enzymes, β and γ -secretases, results in the creation of a 40 or a 42 amino acids long peptide – namely $A\beta$ [11]. Mutations in any one of the secretases leads to increased production of $A\beta$, and a higher susceptibility for Alzheimer's disease [12].

Current structure determination techniques include NMR, X-ray crystallography and cryo-EM (cryoelectron microscopy).

Although X-ray crystallography is the most accurate technique for determining the structure of large proteins, NMR is considered more suitable for elucidating the structure of short peptides at physiological conditions [13]. Several studies have attempted to determine the structure of the $A\beta$ peptide using NMR, but disconcertingly most of them had to use hydrophobic solvents such as TFE (trifluoroethanol) [14–16] and HFIP (hexafluoroisopropanol) [17] to increase solubility. Adding TFE and HFIP to $A\beta$ can induce conformational changes and impose an artificial α -helical structure [18], thus providing the researchers with biased structural information. Remarkably, one attempt to determine the NMR structure without TFE at physiological conditions, revealed $A\beta$ to adopt a 3_{10} -helical structure incongruent with previous α -helical observations [19]. Finally, another NMR study used micelles to determine the structure of $A\beta_{10-35}$ and found it to possess two α -helical regions within residues 13–23 and 30–35 [20].

In this study, we determine the structure of a soluble $A\beta$ peptide corresponding to residues 17–34 which comprises the

Abbreviations: $A\beta$, amyloid β ; AChR, acetylcholine receptor; APP, amyloid precursor protein; HFIP, hexafluoroisopropanol; nAChR, nicotinic acetylcholine; TFE, trifluoroethanol receptor.

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The atomic coordinates of the energy-minimized average coordinates of the $A\beta_{17-34}$ have been deposited in the Protein Data Bank under PDB ID 2MJ1, and the proton chemical shifts and J-coupling data have been deposited in the Biological Magnetic Resonance Data Bank under BMRB ID 19701.

nucleation site of A β using NMR spectroscopy at physiological conditions. We find it to adopt a partial α -helical conformation in agreement with bioinformatic secondary structure predictions. This finding sheds light on the A β conformations in Alzheimer's disease and provides a snapshot of the intermediate structure prior to conversion into amyloid plaques.

EXPERIMENTAL

A β peptides

A β peptides corresponding to residues 17–34 were purchased and purified by the purveyor to a purity level of >99.9%, using HPLC. These peptides contained two glutamate or arginine residues at the N- and C-termini for increased solubility. The full amino acid sequence of the purchased A β was EELVFFAEDVGSNK-GAIGLEE (Peptide 2.0 Inc. Company) and RRLVFFAEDVG-SNKGAIIGLRR (Sigma-Aldrich). The peptide mass was verified using MALDI-TOF mass spectroscopy. To dissolve the peptides, phosphate and acetate aqueous buffers were prepared. None of the buffers contained hydrophobic solvents known to solubilize and stabilize α -helices (i.e. TFE or HFPI) or surfactants known to dissolve hydrophobic peptides (i.e. SDS). Henceforth, the A β _{17–34} peptide refers to the peptide flanked by two glutamates.

NMR sample preparation

To prepare a sample for NMR measurements, 1 mg of A β _{17–34} peptide was dissolved in 300 μ l of 50 mM phosphate buffer (pH 7). To this solution, 15 μ l of D₂O (Sigma-Aldrich) and 1 μ l of sodium azide (100 μ M) (Sigma-Aldrich) were added to a final concentration of 50 mM PO₄, 5% D₂O and 0.3 μ M NaN₃. This NMR sample was pipetted into a Shigemitsu tube (Shigemitsu Corporation) and sealed with a plunger.

NMR experiments

¹H-NMR experiments were performed on a 600 MHz Bruker NMR spectrometer at 278 K. After calibration, 2D TOCSY and ROESY spectra were measured [21–23]) with 2048 and 1024 points in the F2 and F1 dimensions, respectively and 160 scans. The WATERGATE pulse sequence was used in order to suppress the water signal [24,25]. Mixing time for ROESY experiment was set to 400 ms for proper magnetization transfer. Spectral processing was carried out using Bruker TopSpin (version 3.2) software. Spectrum and chemical shift assignment were carried out using the sequential assignment technique developed by Kurt Wüthrich [13].

Structure determination

Structure determination was performed using the Crystallography and NMR system (CNS, version 1.3) software suite [26]. Structures were calculated with the distance geometry and simulated

annealing protocols followed by energy minimization using distance and dihedral angle constraints. Dihedral angle constraints were derived from the ³J_{HNH α} -couplings measured in the TOCSY spectrum processed with 4096 points in the F2 dimension. Distance constraints were calculated from cross-peak height in the ROESY spectrum.

RESULTS

Peptide solubility

To solubilize A β , several approaches were taken. One approach was to add two arginines at the N- and C-termini. This addition increased the theoretical peptide pI from 4.37 (without arginines) to 11.54, but did not improve solubility in acetate and phosphate buffers at pH 1, 7 and 10. Another approach was to add two glutamic acids at the N- and C-termini. This addition decreased the theoretical peptide pI from 4.37 (without glutamates) to 3.83, and improved solubility drastically. The glutamic acid peptides were soluble in both 50 mM acetate and 50 mM phosphate buffers at pH 7 but not in water.

NMR sample preparation

NMR samples containing 1 mg of A β _{17–34} peptide dissolved in 300 μ l of phosphate buffer displayed a pH of 7 (uncorrected for the isotope effect). The NMR samples did not show visible sedimentation or aggregation following 5 min centrifugation at 14000 Rev. per min at 278 K.

NMR measurements

The chemical shifts of all protons of the A β _{17–34} peptide were identified in the TOCSY spectrum using Wüthrich's sequential assignment technique. Figure 1 shows the TOCSY spectrum of this peptide with its 18 amino acid systems. The four glutamic acid residues, two at the C-terminus and two at the N-terminus were not detected due to their high flexibility. The spectrum showed one system per amino acid and so was suggestive of a single conformation.

The ROESY spectrum (not shown) displayed intermediate range H α (*i*)/HN(*i* + 4) cross-peaks between the following amino acid pairs: V18-E22, F20-V24, A21-G25, D23-N27, V24-K28, G25-G29, S26-A30, N27-I31, K28-I32, G29-G33 and A30-L34 (summarized in Figure 2). These cross-peaks were indicative of an α -helix conformation. Additional data supporting the α -helical structure include ³J_{HNH α} -couplings of residues F19, F20, D23, G25 and G33 with values of 6 Hz or less (Figure 2). Also shown in Figure 2 is a summary of the H α (*i*), HN(*i* + 1) connectivities used for the sequential assignment. Interestingly, no HN-H α cross-peak was observed between residues N27 and K28, suggesting a flexible random coil (or kink) in the helix around residues N27 and K28.

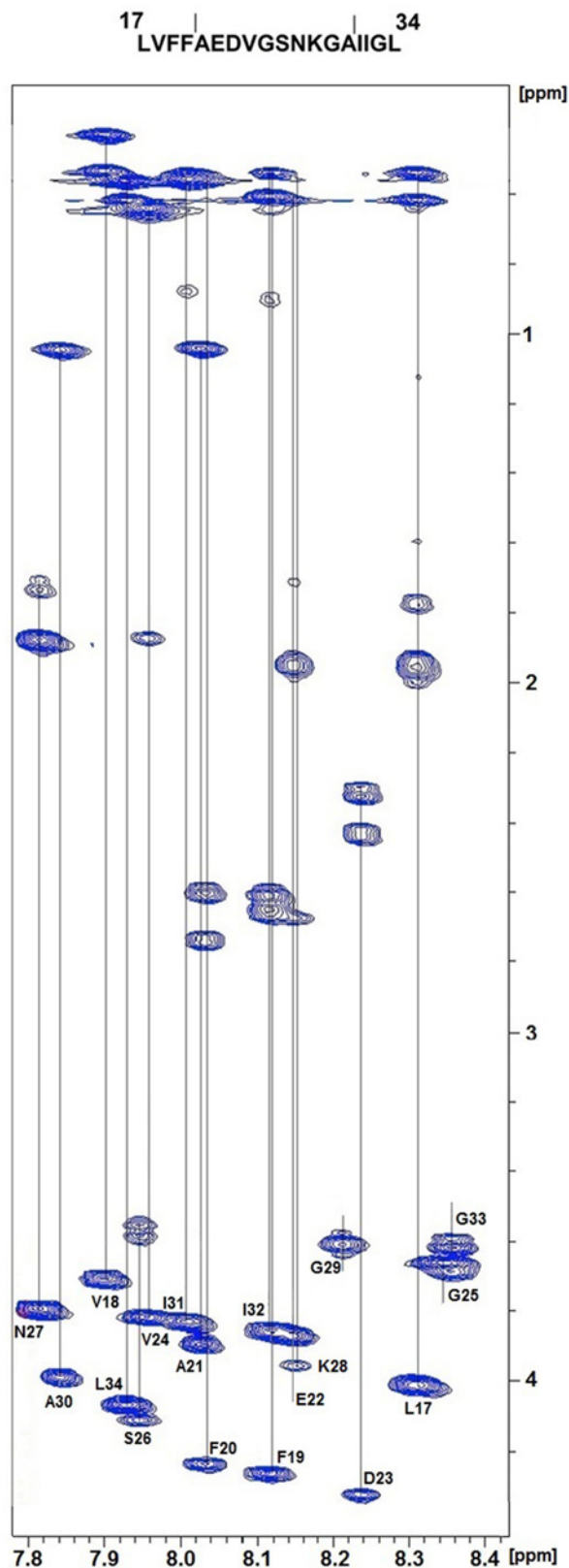


Figure 1 Fingerprint region of TOCSY spectrum of A β ₁₇₋₃₄ peptide. Each line represents one amino acid system.

Table 1 NMR constrains and structural statistics for 10 A β ₁₇₋₃₄ peptide structures

Measurements	
NMR distance constrains	
Total constrains	86
Medium range ($ i-j < 4$)	11
Torsion angle constrains (φ)	5
Mean RMSD values	
Backbone atoms	1.48
Backbone atoms of α -helix	0.28

NMR structure

The lowest energy NMR structures of the soluble A β ₁₇₋₃₄ peptide were found to adopt an α -helix conformation for most of the residues, with a break around S26, N27 and K28 in six out of ten ensemble structures. Figure 3 shows the average structure of the A β ₁₇₋₃₄ peptide. The α -helix spans residues 19–26 and 28–33 with a pronounced kink around residues S26, N27 and K28. Interestingly, the α -helix of residues 23–26 and 28–33 was not previously described. Please note that residues 17–23 and 33–34 did not adopt helical conformation probably due them being terminal residues. Table 1 illustrates the statistical data of the final set of ten structures calculated using CNS. Distance constraints from the ROESY spectrum originated mainly from the fingerprint region. Torsion angle constraints were measured in the TOCSY spectrum with increased processing resolution and identified for residues F19, F20, D23, G25 and G33.

Table 2 shows a Chou–Fasman secondary structure prediction of the A β ₁₋₄₀ peptide. [27]. The secondary structure was also predicted using more modern servers (NetsurfP [28], Jpred [29], I-TASSER [30]); however, the confidence level in these was not high. It is interesting to note that the A β peptide shows a high propensity for both α and β secondary structure. This tendency is an important hallmark in secondary structure conversion of the A β peptide. Our α -helical structure is in agreement with the Chou–Fasman prediction and note that the kink observed around residues S26, and N27 coincides with residues G25 (α -helix propensity of 0.43), S26 (α -helix propensity of 0.57) and N27 (α -helix propensity of 0.76), which possess the lowest α -helix propensities of all amino acids, except proline [31].

DISCUSSION

Several studies have shown that soluble A β and Alzheimer's disease are intertwined and that soluble A β peptide contributes to disease development and progression [32]. Nevertheless, many important details about soluble A β including its structure are missing, biased or contradicting.

Several studies found that A β peptides possess helical or random coil structures under non-physiological conditions. One NMR study found A β ₁₋₄₀ to adopt an 3_{10} -helix in residues 13–23 in aqueous environment (PDB ID 2LFM) [19]. Another NMR

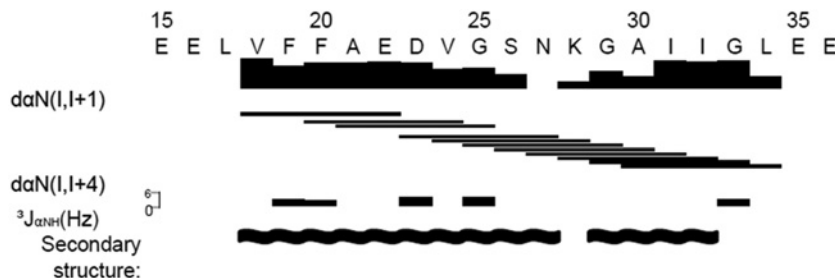


Figure 2 Summary of NMR data used for the sequential assignment and secondary structure determination of $A\beta_{17-34}$ peptide

The data were obtained from the ROESY spectrum recorded at 278 K and pH 7 with 400 ms mixing time. Line thickness indicates the relative cross-peak height of the sequential NOE connectivities.

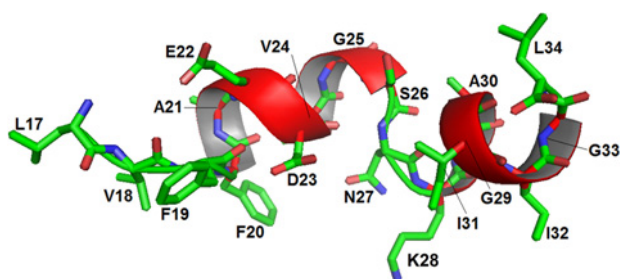


Figure 3 Average NMR structure of the $A\beta_{17-34}$ peptide

Shown is a ribbon diagram and stick representation of the $A\beta_{17-34}$ structure. Notice the two α -helices formed by segments 19–26 and 28–33 with a break around residues S26, N27 and K28.

Table 2 Chou–Fasman secondary structure prediction of $A\beta_{1-40}$

	1	10	20	30	40
Sequence	DAEFRHDSGYEVHHQKLVFFAEDVGSNKGAIIGLMVGGVV				
Helix	<----->		<----->		<----->
Sheet	EEEEEEEEEE				
Turns	T	T	T	T T	T

study found $A\beta_{10-35}$ to adopt an α -helix in residues 13–23 and 30–35 inside SDS micelles [20]. However, at pH 7.3 $A\beta_{10-35}$ is unstructured and adopts a compact random-coil conformation. In another study, the NMR structure of $A\beta_{1-42}$ in a solution containing HFPI/H₂O 30:70 adopts an α -helical conformation in residues 8–25 and 28–38 (PDB ID 1Z0Q) [17]. This finding is similar to ours; however we did not use solubilizing solvents capable of altering protein conformation.

In addition, $A\beta$ was also found to adopt β -hairpin and β -sheet conformations. In one FTIR (Fourier-transform infrared) study, $A\beta$ oligomers were found to adopt an antiparallel β -sheet structure [33]. Another study suggested that $A\beta_{12-42}$ adopts an antiparallel β -sheet conformation with a turn at residues 25–28 [34]. Finally, a third NMR study of $A\beta_{1-40}$ in complex with a phage-display selected antibody, found that residues 17–36 adopt a β -hairpin conformation with a turn composed of residues 23–30 (PDB ID 20TK) [35].

Our findings suggest that the α -helix is longer than previously observed [19] and it is in agreement with secondary structure

predictions (Table 2). We did not observe the α -helix described in residues L17 and V18 (Table 2), probably because the peptide is truncated at residue 17. Also, we did not observe an α -helical conformation for residue L34 because the peptide is truncated here too.

Interaction with nAChR (nicotinic acetylcholine receptor)

Since the first report of $A\beta$ interacting with the nAChR was published [36] several studies attempted to determine the binding mode ([37] and references therein). Before binding the AChR (acetylcholine receptor), $A\beta$ possesses a helical structure with a kink around residue K28. As was reported by us previously [37], K28 is an important anchor point for the amyloid in AChR. A kink around this residue would indeed facilitate anchoring and interaction between the $A\beta$ and AChR. One could hypothesize that binding to AChR condemns it to undergo structural change from helix to β -sheet inside the binding pocket and leaves the peptide in β -sheet form, allowing it to form fibrils and plaques later. In this study, we show the intermediate soluble state of the $A\beta$, which could facilitate binding to the AChR.

Conformational transition

Interestingly, based on secondary structure propensities, the $A\beta$ peptide shows a dual preference for α -helix and for β -sheet (Table 2). This double preference may be attributed to the environment, and $A\beta$ embedded in membranes preferentially adopts an α -helical conformation. Contrarily, in aqueous environment the equilibrium between these states is pushed towards a β -sheet structure, and once this occurs, aggregation and sedimentation follow. In this study, we show the delicate intermediate state of the $A\beta$ peptide solubilized by four glutamate residues. Using NMR, we show that $A\beta$ peptide possesses a stable α -helical structure within residues F20–S26 and K28–G33 with a break around S26–N27–K28. The helical structure was not previously reported in an aqueous environment in the absence of organic solvents, and only a 3_{10} -helix was observed. Another important finding of this research is a possible explanation regarding amyloids structural change from α -helix to β -sheet. Based on our structure, the existence of a kink in the middle of the peptide, which breaks it into two helices allows the peptide to undergo structural changes from

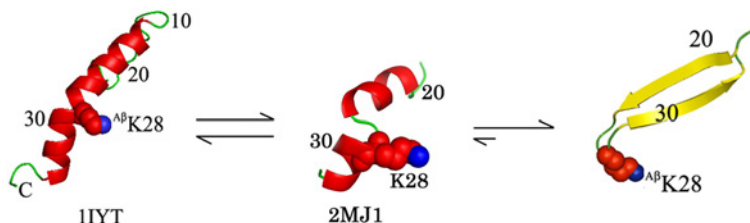


Figure 4 Proposed model of A β structural changes

Shown from left to right is a proposed mechanism of A β transition from α -helix to β -sheet. Shown on the left is the α -helical solution structure of A β _{1–42} in a membrane-like environment (PDB ID 1IYT) [15]. Shown in the middle is our average NMR structure of the A β _{17–34} peptide (PDB ID 2MJ1). Shown on the right is the β -hairpin conformation of the A β peptide [37]).

α -helix to β -sheet once the two halves come close to one another (Figure 4). Where it not for the solubilizing glutamate residues, then A β _{17–34} would aggregate immediately upon leaving the membrane.

Conclusions

This study sheds light on the soluble structure of A β and broadens our biomolecular understanding of Alzheimer's diseases. To the best of our knowledge, this study is the first to report an α -helix in A β in aqueous environment and at physiological conditions. The structure provides an important snapshot of the intermediate state between the pure α -helix and β -sheet conformations, and contributes to our understanding of the conformational transition of A β in Alzheimer's disease.

AUTHOR CONTRIBUTION

Abraham Samson conceived the study and wrote the paper. Genadiy Fonar measured the NMR spectra and calculated the structures.

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