



Cite this: *RSC Adv.*, 2018, 8, 41705

Atomistic investigation of an Iowa Amyloid- β trimer in aqueous solution†

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The self-assembly of Amyloid beta ($A\beta$) peptides are widely accepted to associate with Alzheimer's disease (AD) *via* several proposed mechanisms. Because $A\beta$ oligomers exist in a complicated environment consisting of various forms of $A\beta$, including oligomers, protofibrils, and fibrils, their structure has not been well understood. The negatively charged residue D23 is one of the critical residues of the $A\beta$ peptide as it is located in the central hydrophobic domain of the $A\beta$ N-terminal and forms a salt-bridge D23-K28, which helps stabilize the loop domain. In the familial Iowa (D23N) mutant, the total net charge of $A\beta$ oligomers decreases, resulting in the decrease of electrostatic repulsion between D23N $A\beta$ monomers and thus the increase in their self-aggregation rate. In this work, the impact of the D23N mutation on $3A\beta_{11-40}$ trimer was characterized utilizing temperature replica exchange molecular dynamics (REMD) simulations. Our simulation reveals that D23N mutation significantly enhances the affinity between the constituting chains in the trimer, increases the β -content (especially in the sequence 21–23), and shifts the β -strand hydrophobic core from crossing arrangement to parallel arrangement, which is consistent with the increase in self-aggregation rate. Molecular docking indicates that the $A\beta$ fibril-binding ligands bind to the D23N and WT forms at different poses. These compounds prefer to bind to the N-terminal β -strand of the D23N mutant trimer, while they mostly bind to the N-terminal loop region of the WT. It is important to take into account the difference in the binding of ligands to mutant and wild type $A\beta$ peptides in designing efficient inhibitors for various types of AD.

Received 13th September 2018
Accepted 4th December 2018

DOI: 10.1039/c8ra07615d

rsc.li/rsc-advances

Introduction

World Health Organization (WHO) reported that dementia affected approximately 50 million people worldwide in 2017 and the number of new patients could increase to *ca.* 10 million annually.¹ Alzheimer's disease (AD) is the most common type of dementia (60–70% cases),¹ which is strongly linked to the self-aggregation of the Amyloid beta ($A\beta$) peptides.² $A\beta$ peptides consist of 36–43 residues formed *via* the proteolysis of the transmembrane Amyloid Precursor Protein (APP). According to the Amyloid cascade hypothesis,^{3–5} $A\beta$ peptides self-assemble into $A\beta$ oligomers, which have been shown to be neurotoxic *via* several mechanisms, including damaging neurites and synapses^{2,6} or forming transmembrane

pores that disrupt Ca^{2+} homeostasis.^{7,8} The most abundant form of $A\beta$ peptides consists of 40 amino acids ($A\beta_{40}$) and accounts for approximately 80–90% of the total population.⁹ The development of AD therapy relies on the understanding of $A\beta$ oligomer structures,^{10–13} unfortunately, because $A\beta$ oligomers exist in complicated environments comprising of numerous forms of oligomers and fibrils, their structure has not been well characterized.^{11,14} Therefore, molecular dynamics simulation has been one of the main tools in assessing the structure of $A\beta$ oligomers.^{15–17}

The self-assembly process of $A\beta$ peptides highly depends on their amino acid sequence.¹⁸ Mutations can alter the $A\beta$ oligomer structures and their properties. Especially, the common mutations in the central hydrophobic domain of the $A\beta$ N-terminal, including Flemish (A21G),¹⁹ Dutch (E22Q),²⁰ Italian (E22K),²¹ Arctic (E22G),²² and Iowa (D23N)²³ significantly affect the dynamics and conformations of the $A\beta$ oligomers and fibrils. It is known that the central hydrophobic domains of an $A\beta$ chain forms rigid β -structure that in turn forms strong interaction with other $A\beta$ chains, leading to $A\beta$ self-aggregation. Inhibitors that target the N-terminal β -strand have been shown to successfully inhibit $A\beta$ self-aggregation.^{24–26}

In the N-terminal hydrophobic domain, D23 stabilizes the loop region of $A\beta$ peptides by forming a salt-bridge to the

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† Electronic supplementary information (ESI) available: List of temperatures in REMD simulations. Additional figures displaying the diffusion of the first replica in the entire temperature space; the secondary structure distribution per residue of D23N $3A\beta_{11-40}$ in different simulation intervals; the RMSD and FEL of D23N and WT $A\beta_{11-40}$ monomers; and docked conformations of ligands to monomers. See DOI: 10.1039/c8ra07615d



residue K28 and polar contacts to other residues in loop domain.^{27–29} Substituting the negatively charged residue D23 by another residue could significantly alter the structure of A β peptides. In the familial Iowa mutant, D23 is replaced by the neutral residue asparagine,²³ which reduces the total net charge and thus the electrostatic repulsion between constituting chains of A β oligomers. Consequently, the self-aggregation rate of D23N oligomers has been shown to increase.³⁰ Therefore, studying the impact of the D23N mutation on the self-assembly has been carried out for a number of A β peptide systems, including A β_{21-30} ,³¹ A β_{10-35} ,³² A β_{1-40} ,³³ A β_{1-42} ,³⁴ 2A β_{1-40} ,³⁵ and 6A β_{16-35} .³⁶ However, the effect of D23N mutation on A β_{40} trimer (3A β_{40}), one of the most neurotoxic A β oligomers,^{10,11} has not been revealed.

We are interested in A β_{40} as it is the most abundance form of A β . Although A β_{42} oligomers are more neurotoxic than A β_{40} oligomers, the population of A β_{40} peptides is approximately ten times higher than that of A β_{42} .⁹ Thus, structural investigation of A β_{40} and A β_{42} are both important. In addition, structure–neurotoxicity relationship has been established for A β_{40} oligomers but not for A β_{42} oligomers.³⁷ In order to gain insights into the changes in the structure and dynamics of the neurotoxic A β trimer in aqueous environment upon D23N mutation, we investigated the D23N 3A β_{11-40} mutant in comparison with the wild type (WT) trimer using extensive replica exchange molecular dynamic (REMD) simulation over 20 000 ns in total. It is noted that the D23N A β peptides are able to form both parallel and anti-parallel motifs;^{38,39} however, in this work, we only considered the D23N 3A β_{11-40} starting from the parallel structure. Our results indicate that the D23N mutant has higher β -structure and stronger inter-chain interaction. We also identified seven optimized conformations of solvated D23N 3A β_{11-40} using the combination of free energy landscape and clustering methods. Finally, molecular docking method was used to evaluate the binding affinity of available inhibitors to D23N and WT A β_{11-40} trimer and monomer in order to reveal the influence of D23N mutant on AD therapy.

Materials and methods

Initial conformation

The three-dimensional fibril-like conformation of WT 3A β_{11-40} peptide was taken from the two-fold 12A β_{11-40} fibril.⁴⁰ Then, the residue D23 was mutated to N using the mutagenesis tool in Pymol (Fig. 1).²⁹ The backbone of trimer was not changed, but the sidechain of D23 was slightly different to N23. The distances between N23 and neighboring atoms remained larger than the corresponding atomic van der Waals (vdW) radii. D23N 3A β_{11-40} was then inserted into a dodecahedron periodic boundary condition (PBC) box with the volume of ~ 463 nm³ and solvated with 12 852 water molecules. The mutant trimer was treated with AMBER99SB-ILDN force field⁴¹ and water molecules were presented utilizing TIP3P water model⁴² as described in previous works.^{29,43}

In addition, the D23N and WT A β_{11-40} monomers were derived from the trimer systems. Both monomers were solvated

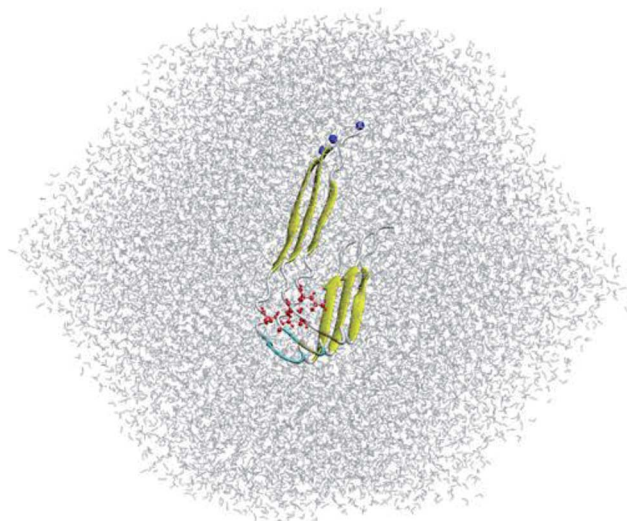


Fig. 1 Input conformation of solvated D23N 3A β_{11-40} . N23 residues are highlighted in red. N-termini are indicated with blue balls.

in a dodecahedron PBC box with size of ~ 270 nm³ using ~ 8600 water molecules. Na⁺ ions were added to WT A β_{11-40} monomer in order to neutralize the system.

Atomistic simulations

Atomistic simulations were carried out as previously described.^{29,44} GROMACS 5.0.6 (ref. 45) was employed to simulate the solvated D23N 3A β_{11-40} system. The non-covalent bond pair cut-off was set at 0.9 nm. The electrostatic potential was treated using the fast smooth particle-mesh Ewald electrostatic method.⁴⁶ The vdW interaction was set at 0.9 nm. The first step of simulation was energy minimization with the steepest descent method, which was followed by a 500 ps positionally restraint simulation in NVT ensemble. REMD simulation was then applied to sample the conformations of solvated D23N

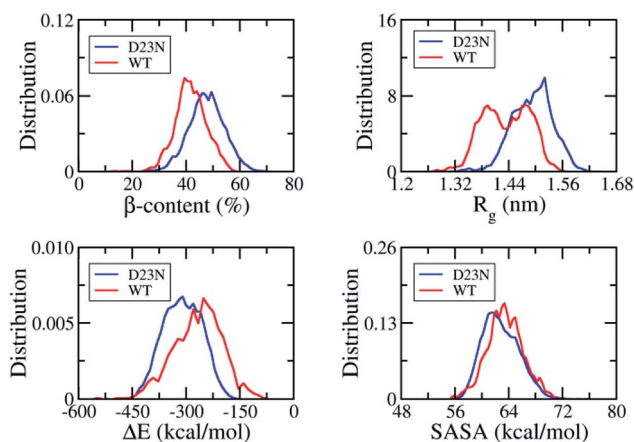


Fig. 2 Distributions of D23N 3A β_{11-40} (red) and WT 3A β_{11-40} (blue) in REMD simulation time interval of 300–417 ns at 300 K. The values of WT 3A β_{11-40} were reproduced with permission of Royal Chemistry Society.²⁹



$3A\beta_{11-40}$. 48 replicas at various temperatures ranging from 295 to 382 K (ESI†) were simulated as reported in our previous works.^{29,43} During REMD simulation, the exchanges were performed every 1 ps. Each replica was simulated over 417 ns. The solvated D23N and WT $A\beta_{11-40}$ monomers were also simulated for 100 ns at 300 K in order to generate the target for molecular docking. The trimer conformations in REMD simulations were recorded every 10 ps, while monomeric conformations in MD simulations were monitored every 1 ps.

Structural analysis

The Defined Secondary Structure of Protein (DSSP) method was employed to predict the secondary structure term of the D23N $3A\beta_{11-40}$.⁴⁷ The inter-/intra-molecular sidechain (inter-/intra-SC) contacts were counted when the spacing between non-hydrogen atoms are smaller than 0.45 nm. The inter-/intra-molecular hydrogen bond (HB) contacts were counted when the spacing between donor and acceptor is smaller than 0.35 nm and the angle between acceptor–hydrogen–donor is larger than 135°. The collective variable free energy landscape (FEL) was generated with two reaction coordinates, including the C_α root-mean-square deviation (RMSD) and gyration of radius R_g . The non-covalent bond interaction energy and solvent surface access area (SASA) were calculated using GROMACS tools.⁴⁵ The collision cross section (CCS) was computed using the Ion Mobility Projection Approximation Calculation Tool (IMPACT).⁴⁸

Binding free energy calculations

The binding free energy of an $A\beta$ peptide chain to the other chains within D23N $3A\beta_{11-40}$ was calculated using MM/PBSA method^{49–51} as previously described.²⁹

Molecular docking

The binding configuration and affinity of available inhibitors to $A\beta$ trimers and monomers were investigated using Autodock Vina package⁵² as described previously.⁵³ The input file of docking was prepared using AutodockTools 1.5.6.⁵⁴ The docking grid was chosen at the center of mass of the peptide with the size of 60 × 60 × 60 nm, which is large enough to cover the entire peptides. The exhaustiveness was set as 400.

Results and discussion

REMD simulations of solvated D23N $3A\beta_{11-40}$

REMD simulation is known to be one of the most efficient sampling methods to evaluate the self-aggregation of $A\beta$ peptides.^{15,16,55,56} Because simulation of the self-aggregation process of $A\beta$ peptides is very slow and it is highly unlikely to reach the stable structure of $A\beta$ oligomers starting from a random structure, input conformations were obtained from random fibril-like structures,^{29,43,44} which was also applied in the case of $A\beta_{11-40}$ trimer and its D23N mutant. REMD simulations were sufficient as the mean exchange rate of *ca.* 32% and the $A\beta$ conformations entirely walked through the temperature space (Fig. S1†). The first 300 ns of REMD simulations were omitted from any analysis due to sidestepping any inclination toward

the starting conformation. All computational metrics of the solvated D23N $3A\beta_{11-40}$ were only evaluated over the last 117 ns of REMD simulation at 300 K.

The simulation reached the equilibrium state at 300 K after 300 ns as indicated by the superposition of secondary structure metrics over various intervals 300–417 and 300–360 ns (Fig. S2†). All metrics were computed over all the snapshots of the simulation window 300–417 ns. As shown in Fig. 2, the β -content value diffuses in the range of 11–71% with the mean value of $\sim 48 \pm 7\%$. It is significantly larger than that of WT $3A\beta_{11-40}$ ($\sim 42 \pm 6\%$).²⁹ The shift of β -content distribution under the effect of D23N mutation is clearly shown in Fig. 2. R_g fluctuates in the range of 1.28–1.64 nm with an averaged value of $\sim 1.49 \pm 0.05$ nm, which is larger than the corresponding value of the WT trimer ($\sim 1.43 \pm 0.05$ nm).²⁹ SASA of D23N $3A\beta_{11-40}$ ($\sim 62.84 \pm 2.66$ nm²) is slightly smaller than that of WT $3A\beta_{11-40}$ ($\sim 63.51 \pm 2.59$ nm²).²⁹

The non-covalent bond interaction energy between peptide chains of D23N $3A\beta_{11-40}$ was obtained using the GROMACS tool 4.3, which falls in the range from -515.7 to -160.5 kcal mol⁻¹ (Fig. 2). The average interaction energy is -310.2 ± 53.3 kcal mol⁻¹, which is significantly larger than that of WT $3A\beta_{11-40}$ (-267.0 ± 67.2 kcal mol⁻¹).²⁹ The stronger interaction energy between establishing chains is observed when the electrostatic repulsion is reduced. The obtained results indicated that the sidechain charge–charge interactions contribute to the instability of the WT trimer.⁵⁷ This result implies that D23N $3A\beta_{11-40}$ is more stable and forms at higher rate in comparison to the WT trimer.

Effects of D23N mutation on the distribution of secondary structure parameters per residue

Fig. 3 shows the distribution of secondary structure terms per residues of the D23N $3A\beta_{11-40}$ including β -, α -, turn-, and coil-structures. Obtained results are consistent with the available solid state NMR data that showed a high neurotoxic $A\beta_{40}$ oligomer possessing stable N-terminal β -strands.⁵⁸ D23N $3A\beta_{11-40}$ adopts parallel U-shape structure consisting of five domains, which is in good agreement with a recent study.³⁸ The sequences 11–14, 24–30, and 36–40 mostly form coil structure, while β -structure is dominant in the sequences 15–23 and 31–35. Residues 21–23 in the D23N mutant possess remarkably higher β -structure content compared to those in the WT trimer. Moreover, β -structure content is also observed to increase in other residues, including the sequence 12–20, while it only decreases in several residues, including the sequence 31–34. The loop region (sequence 24–28) forms more coil-structure and less α /turn-structure in comparison with the WT form. Overall, the significant decrease in α -structure, an intermediate structure in the $A\beta$ folding progress,^{59,60} together with the significant increase in β -structure could speed up the self-aggregation of D23N $3A\beta$ in comparison with that of the WT trimer.

Non-covalent bond contacts

The stability of $A\beta$ oligomer depends on the intermolecular non-covalent bonds between constituting chains, involving



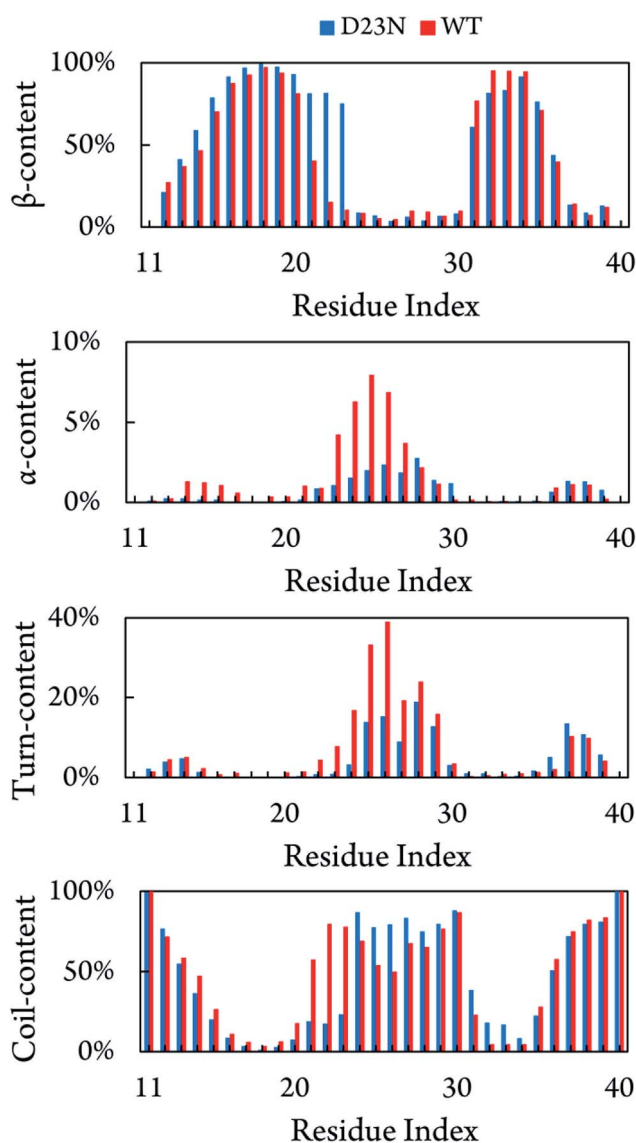


Fig. 3 Secondary structure terms per residues averaged for all three chains of D23N $3A\beta_{11-40}$ and WT $3A\beta_{11-40}$.²⁹ The values of WT $3A\beta_{11-40}$ were reproduced with permission of Royal Chemistry Society.²⁹

inter-SC and inter-HB contacts. The distributions of these values are shown in Fig. 4. As mentioned above, D23N $A\beta_{11-40}$ forms more inter-SC contact (66.0 ± 4.5) than the WT system (61.8 ± 5.2).²⁹ Similarly, D23N $A\beta_{11-40}$ adopts more inter-HB contact (12.1 ± 5.0) than WT trimer (10.2 ± 1.7).²⁹ Moreover, the number of intramolecular contacts of the D23N $3A\beta_{11-40}$ (135.7 ± 7.8) is significantly smaller than those of WT $3A\beta_{11-40}$ (171.2 ± 8.3).²⁹ Likewise, the intra-HB of D23N $3A\beta_{11-40}$ (6.8 ± 2.0) is lower than that of the WT trimer (7.6 ± 2.9).²⁹ This result is consistent with the significant increase in interaction energy between the peptide chains of the trimers obtained using the GROMACS tools described above, as well as with the increase in Gibbs binding free energy between peptide chains derived from MM/PBSA method (*vide infra*).

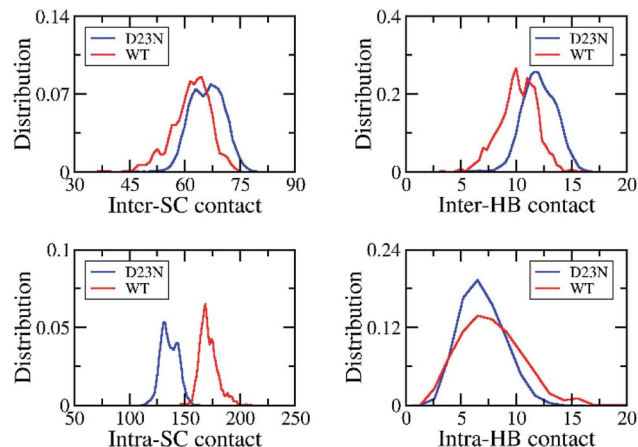


Fig. 4 The distribution of inter/intra-molecular SC and HB contacts of the solvated D23 and WT $3A\beta_{11-40}$. The results of WT system was reproduced with permission of Royal of Chemistry Society.²⁹

Free energy landscape

The collective variable free energy landscape (FEL) of D23N $3A\beta_{11-40}$ was generated using the GROMACS tool named “sham”.⁴⁵ C_{α} RMSD and R_g were chosen as the coordinates as they were successfully used to obtain the FEL for other $A\beta$ peptide systems.⁶¹⁻⁶³ The FEL of solvated D23N $3A\beta_{11-40}$ is displayed in Fig. 5, which contains four minima centered at (RMSD; R_g) coordinates of (0.48; 1.52), (1.50; 0.61), (0.68; 1.46), (0.66; 1.48). Applying clustering method on the refined MD snapshots located in the free energy holes, we found seven representative conformations of D23N $3A\beta_{11-40}$. These conformations are noted as A, B, C, D, E, F, and G as shown in Fig. 5.

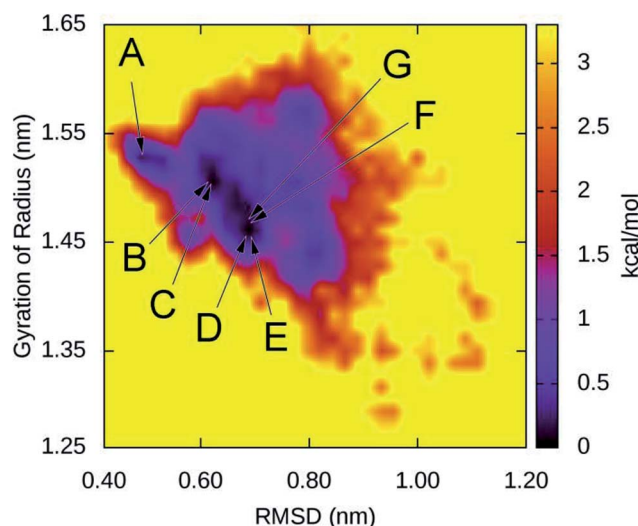


Fig. 5 Collective variable free energy landscape of D23N $3A\beta_{11-40}$ generated with the reaction coordinates of C_{α} RMSD and R_g . Seven minima, namely A, B, C, D, E, F, and G, are found in four free energy holes. The corresponding (RMSD; R_g) coordinates of minima are (0.48; 1.52) for A, (0.61; 1.50) for B and C, (0.68; 0.68) for D and E, and (0.66; 1.48) for F and G.



Table 1 Structural details of the solvated D23N 3A β_{11-40} peptide

	R_g (nm)	SASA (nm ²)	CCS (nm ²)	β (%)
A	1.52	61.70	13.19	51
B	1.51	61.79	13.07	54
C	1.50	63.08	13.38	47
D	1.46	60.46	13.14	48
E	1.46	58.97	12.83	46
F	1.48	63.13	13.54	42
G	1.48	58.80	12.91	58

The selected structural parameters of these conformations are provided in Table 1. The dimensions of these conformations, which are indicated by R_g , SASA, and CCS, are roughly comparable to those found for the representative structures of the WT trimers. The β -content of these structures is greater than 42%, which is significantly higher than that found for the representative structures of WT 3A β , namely MA (37%), MB (40%), and MC (32%).²⁹ These results are consistent with the analysis of all equilibrated snapshots described above.

The 3-D structures of A–G and their populations are shown in Fig. 6. All conformations exhibit the U-shaped parallel structures. As mentioned above, the β -strand of the N-terminal in the D23N mutant forms more β -structure as previously observed by solid state NMR.⁵⁸ It is worthy to note that the central hydrophobic domain is solvent accessible, to which other A β peptides could dock, resulting in the self-aggregation.⁶⁴ Unlike the β -strands in the WT trimer that form crossing structure,²⁹ the β -strands in D23N 3A β_{11-40} are parallel to each other. Docking of additional monomer to the parallel structure of D23N 3A β_{11-40} is much more efficient than that to the crossing structure of the WT trimer.⁶⁴

Binding affinity between peptide chains in A β_{11-40} trimers

The increase in the interaction energy and non-covalent bond contacts upon D23N mutation of 3A β_{11-40} described above suggest that the peptide chains in the mutant trimer bind more strongly to each other than in the WT trimer. To further access this result, we performed MM/PBSA calculation to evaluate the Gibbs free binding energy (ΔG_{bind}) of a peptide chain to the other chains in D23N 3A β_{11-40} , using the same protocol previously described for WT 3A β_{11-40} .²⁹ Evaluating binding affinity between peptide chains of the trimer using ΔG_{bind} obtained with MM/PBSA method is more accurate than using the interaction energy obtained with GROMACS tool. The parameters obtained with MM/PBSA method are provided in Table 2. Replacing the negatively-charged D23 residue by a neutral N residue results in remarkably stronger electrostatic interaction (ΔE_{elec}) and slightly stronger vdW interaction (ΔE_{vdw}) as a result of weaker repulsion force between the peptide chains. The polar interaction is significantly stronger in the mutant, which off-set the increase in affinity between the chains. Nevertheless, the magnitude of ΔG_{bind} increases significantly (by $\sim 22\%$) when D23 is mutated to N, which is consistent with the increase in the magnitude of interaction energy obtained with GROMACS tool.

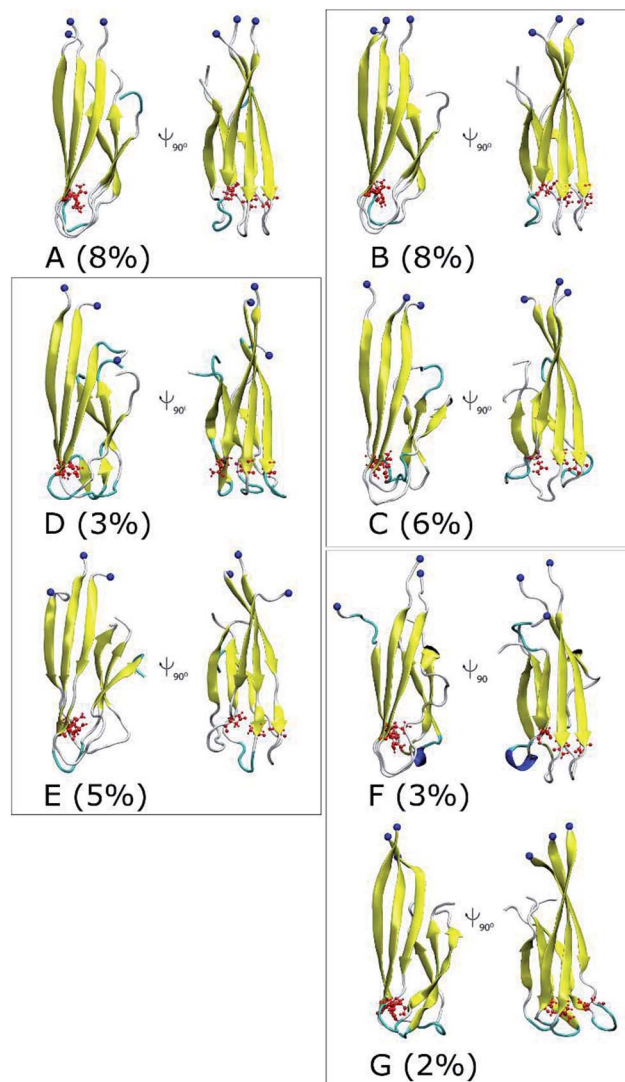


Fig. 6 Representative structures of the solvated D23N 3A β_{11-40} obtained from the combination of FEL and clustering methods. The blue balls note the N-terminus of D23N trimer.

Binding of inhibitors to A β trimers and monomers

In order to gain more insights into how D23N mutation affects the structure of A β trimers, we compared the binding of twelve experimentally characterized amyloid plaque-binding compounds^{65–71} to D23N 3A β_{11-40} and WT 3A β_{11-40} using Auto-dock Vina.⁵² This tool has been used to study the binding of ligands to amyloid peptides and provided good correlation

Table 2 MMPBSA calculation parameters (kcal mol⁻¹)^a

	ΔE_{elec}	ΔE_{vdw}	ΔG_{sur}	ΔG_{polar}	$-T\Delta S$	ΔG_{bind}
WT ²⁹	-90.08	-144.92	-21.26	153.62	66.82	-35.82 \pm 7.89
D23N	-180.72	-126.39	-24.6	217.78	70.29	-43.67 \pm 15.90

^a ΔE_{elec} = electrostatic interaction energy; ΔE_{vdw} = van der Waals interaction energy; ΔG_{sur} = non-polar free energy of binding; ΔG_{polar} = polar free energy of binding; $-T\Delta S$ = contribution of entropy to Gibbs free energy; the results of the WT system were reproduced with permission of Royal of Chemistry Society.²⁹



Table 3 Binding affinity (kcal mol⁻¹) of selected inhibitors to D23N and WT 3Aβ₁₁₋₄₀

No.	PubChem ID	Ref.	D23N 3Aβ ₁₁₋₄₀	WT 3Aβ ₁₁₋₄₀
1	156 391	65	-5.8	-5.4
2	234 475	66	-5.4	-5.2
3	1 358 096	67	-6.0	-5.7
4	9 835 167	66	-5.7	-5.5
5	9 837 643	66	-5.6	-5.4
6	9 839 907	68	-5.7	-5.5
7	10 133 297	69	-5.7	-5.5
8	10 335 228	66	-5.5	-5.2
9	10 422 457	66	-5.7	-5.5
10	10 444 413	66	-5.7	-5.5
11	10 844 298	70	-6.2	-6.1
12	16 087 303	71	-4.8	-4.9

between predicted and experimental binding affinities.^{72,73} Molecular docking of these ligands was carried out for all representative structures of D23N and WT 3Aβ₁₁₋₄₀. For each system, the binding affinity was averaged from the values obtained for all representative conformations to compensate for the lack of receptor-inhibitor dynamics in Autodock Vina. The exhaustiveness was chosen as 400 to enhance the docking accuracy.

The docking results are presented in Tables 3 and S1†. Except for one ligand, the estimated binding affinity of the ligands to D23N 3Aβ₁₁₋₄₀ is slightly higher than that to WT

3Aβ₁₁₋₄₀, however, the difference is negligible. Fig. 7 shows the binding poses of the ligands to the representative conformations of D23N 3Aβ₁₁₋₄₀ (A–G) and WT 3Aβ₁₁₋₄₀ (MA–MC). Overall, the ligands bind to D23N 3Aβ₁₁₋₄₀ in different positions compared to WT 3Aβ₁₁₋₄₀. The binding poses of the ligands to D23N 3Aβ₁₁₋₄₀ are relatively wide spread, but in most cases they bind to the N-terminal β-strands. While the ligands also bind to the N-terminal domain of WT 3Aβ₁₁₋₄₀, they mostly dock to the loop regions in similar poses in each conformation. This result suggests that these ligands could inhibit the aggregation of D23N and WT Aβ by different mechanisms. Binding of the ligands to the β-strands may inhibit the docking of new Aβ chain to the existing oligomers, while binding of the ligand to the loop regions could lead to the destabilization of the oligomers.

In addition, the inhibitors were also docked to the D23N and WT Aβ₁₁₋₄₀ the representative structures of the monomers obtained from FEL and clustering methods (Fig. S4†). D23N mutation cause significant changes in the structure of Aβ₁₁₋₄₀ monomer (Fig. S4†). The binding poses of the inhibitors to D23N monomer are different from those to WT Aβ₁₁₋₄₀ monomer (Fig. S5†). The average ligand binding affinity of D23N monomer (-5.0 ± 0.2 kcal mol⁻¹) is 0.7 kcal mol⁻¹ smaller than that of D23N trimer (-5.7 ± 0.3 kcal mol⁻¹) (Table S2†). Similarly, the average ligand binding affinity of WT Aβ₁₁₋₄₀ (-4.8 ± 0.2) is 0.7 kcal mol⁻¹ smaller than that of WT 3Aβ₁₁₋₄₀ ($-5.5 \pm$

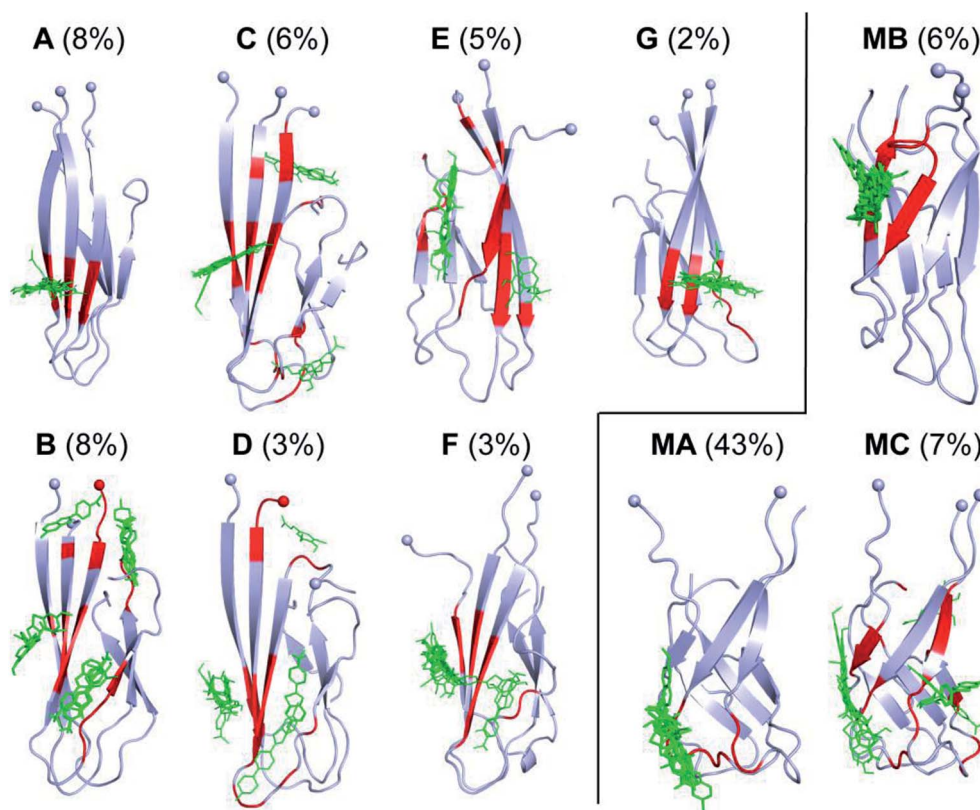


Fig. 7 Binding poses of twelve ligands to the representative conformations of D23N 3Aβ₁₁₋₄₀ (A–G) (this work) and WT 3Aβ₁₁₋₄₀ (MA–MC) (obtained from our previous study)²⁹ generated using Autodock Vina protocol.⁵² The residues highlighted by red color form non-bonded contacts to the ligands. Spheres denote the C_α atoms of the N-terminal.



0.3 kcal mol⁻¹). This result indicates that the inhibitors have higher affinity for the trimers than for the monomers. It also shows that D23N mutation also causes changes in ligand binding poses of the monomers, further corroborating the results obtained for the trimers.

Conclusions

Atomistic structural and dynamic information of the (11–40) truncated Iowa A β mutant trimer (D23N 3A β _{11–40}) were extensively simulated using REMD with 48 replicas over simulation time of 417 ns per replica (~20 000 ns of total simulations time). Replacing the negatively-charged D23 residue by the neutral residue N increases β -structure content from ~42% to ~48%, which occurs concomitantly with a shift in the hydrophobic β -strand core from crossing to parallel arrangement (Fig. 7). Subsequent docking of other A β chains *via* dock-lock mechanism^{64,74} to the existing oligomer with parallel hydrophobic core could occur much more efficiently than to the oligomer with crossing core. The number of non-covalent contacts increases sizably in the D23N mutant trimer, which is consistent with the increase in the magnitude of both the interaction energy (from -267.0 ± 67.2 kcal mol⁻¹ to -310.2 ± 53.3 kcal mol⁻¹) derived with the GROMACS tool and the Gibbs free binding energy (from -35.82 ± 7.89 to -43.67 ± 15.90) obtained with MM/PBSA method. Altogether, our results show that D23N mutation induces significant changes in structure and dynamics of 3A β _{11–40} that allow the mutant peptide to aggregate at faster rate, consistent with previous experimental results on the formation of Iowa A β fibrils.³⁰

Molecular docking shows that D23N mutation causes significant changes of A β _{11–40} structure. These changes lead to differences in ligand binding poses of D23N and WT trimers despite having similar binding affinity. The ligands mostly bind to the N-terminal β -strand of the D23N mutant trimer, while they prefer to bind to the N-terminal loop region of the WT trimer. Likewise, these compounds bind to D23N and WT A β _{11–40} monomers at different binding positions while having the same range in binding affinity. This result provides important information for further study in drug design for different types of AD. Rigorous MD simulations are needed to gain further insights into this aspects.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This research is funded by Vietnam National Foundation for Science and Technology Development (NAFOSTED) under grant number 103.01-2016.48.

Notes and references

1 WHO, Dementia, <http://www.who.int/en/news-room/fact-sheets/detail/dementia>, (accessed July 29, 2018).

- D. J. Selkoe, *Science*, 2002, **298**, 789–791.
- K. Beyreuther and C. L. Masters, *Brain Pathol.*, 1991, **1**, 241–251.
- J. Hardy and D. Allsop, *Trends Pharmacol. Sci.*, 1991, **12**, 383–388.
- D. J. Selkoe, *Neuron*, 1991, **6**, 487–498.
- S. Chimon, M. A. Shaibat, C. R. Jones, D. C. Calero, B. Aizezi and Y. Ishii, *Nat. Struct. Mol. Biol.*, 2007, **14**, 1157.
- T. L. Williams and L. C. Serpell, *FEBS J.*, 2011, **278**, 3905–3917.
- A. Quist, I. Doudevski, H. Lin, R. Azimova, D. Ng, B. Frangione, B. Kagan, J. Ghiso and R. Lal, *Proc. Natl. Acad. Sci. U. S. A.*, 2005, **102**, 10427–10432.
- M. P. Murphy and H. LeVine, *J. Alzheimers Dis.*, 2010, **19**, 311.
- S. J. C. Lee, E. Nam, H. J. Lee, M. G. Savelieff and M. H. Lim, *Chem. Soc. Rev.*, 2017, **46**, 310–323.
- M. K. Jana, R. Cappai, C. L. L. Pham and G. D. Ciccotosto, *J. Neurochem.*, 2016, **136**, 594–608.
- S. T. Ngo, S.-T. Fang, S.-H. Huang, C.-L. Chou, P. D. Q. Huy, M. S. Li and Y.-C. Chen, *J. Chem. Inf. Model.*, 2016, **56**, 1344–1356.
- D. J. Selkoe and J. Hardy, *EMBO Mol. Med.*, 2016, **8**, 595–608.
- S. Banerjee, Z. Sun, E. Y. Hayden, D. B. Teplow and Y. L. Lyubchenko, *ACS Nano*, 2017, **11**, 12202–12209.
- P. H. Nguyen, F. Sterpone, J. M. Campanera, J. Nasica-Labouze and P. Derreumaux, *ACS Chem. Neurosci.*, 2016, **7**, 823–832.
- S. T. Ngo, M. T. Nguyen, N. T. Nguyen and V. V. Vu, *J. Phys. Chem. B*, 2017, **121**, 8467–8474.
- S. T. Ngo, *Commun. Phys.*, 2018, **28**, 265–276.
- L. K. Thompson, *Proc. Natl. Acad. Sci. U. S. A.*, 2003, **100**, 383–385.
- L. Hendriks, C. M. van Duijn, P. Cras, M. Cruts, W. Van Hul, F. van Harskamp, A. Warren, M. G. McInnis, S. E. Antonarakis, J.-J. Martin, A. Hofman and C. Van Broeckhoven, *Nat. Genet.*, 1992, **1**, 218–221.
- E. Levy, M. Carman, I. Fernandez-Madrid, M. Power, I. Lieberburg, S. van Duinen, G. Bots, W. Luyendijk and B. Frangione, *Science*, 1990, **248**, 1124–1126.
- O. Bugiani, A. Padovani, M. Magoni, G. Andora, M. Sgarzi, M. Savoirdo, A. Bizzi, G. Giaccone, G. Rossi and F. Tagliavini, *Neurobiol. Aging*, 1998, **19**, S238.
- C. Nilsberth, A. Westlind-Danielsson, C. B. Eckman, M. M. Condron, K. Axelman, C. Forsell, C. Stenh, J. Luthman, D. B. Teplow, S. G. Younkin, J. Naslund and L. Lannfelt, *Nat. Neurosci.*, 2001, **4**, 887–893.
- T. J. Grabowski, H. S. Cho, J. P. G. Vonsattel, G. W. Rebeck and S. M. Greenberg, *Ann. Neurol.*, 2001, **49**, 697–705.
- D. Sehlin, H. Englund, B. Simu, M. Karlsson, M. Ingelsson, F. Nikolajeff, L. Lannfelt and F. E. Pettersson, *PLoS One*, 2012, **7**, e32014.
- L. Lannfelt, C. Möller, H. Basun, G. Osswald, D. Sehlin, A. Satlin, V. Logovinsky and P. Gellerfors, *Alzheimer's Res. Ther.*, 2014, **6**, 16.
- C. Swanson, *A Study to Evaluate Safety, Tolerability, and Efficacy of BAN2401 in Subjects With Early Alzheimer's Disease*, <https://clinicaltrials.gov/ct2/show/NCT01767311>, (accessed July 29, 2018).



- 27 B. Tarus, J. E. Straub and D. Thirumalai, *J. Mol. Biol.*, 2005, **345**, 1141–1156.
- 28 B. Ma and R. Nussinov, *Proc. Natl. Acad. Sci. U. S. A.*, 2002, **99**, 14126–14131.
- 29 S. T. Ngo, H. M. Hung, D. T. Truong and M. T. Nguyen, *Phys. Chem. Chem. Phys.*, 2017, **19**, 1909–1919.
- 30 X. Yang, G. Meisl, B. Frohm, E. Thulin, T. P. J. Knowles and S. Linse, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, **115**, E5849–E5858.
- 31 B. Tarus, J. E. Straub and D. Thirumalai, *J. Mol. Biol.*, 2008, **379**, 815–829.
- 32 W. Han and Y.-D. Wu, *J. Am. Chem. Soc.*, 2005, **127**, 15408–15416.
- 33 S. Côté, P. Derreumaux and N. Mousseau, *J. Chem. Theory Comput.*, 2011, **7**, 2584–2592.
- 34 S.-H. Chong, J. Yim and S. Ham, *Mol. BioSyst.*, 2013, **9**, 997–1003.
- 35 S. Côté, R. Laghaei, P. Derreumaux and N. Mousseau, *J. Phys. Chem. B*, 2012, **116**, 4043–4055.
- 36 W. Han and Y.-D. Wu, *Proteins*, 2007, **66**, 575–587.
- 37 K. Ono, M. M. Condrón and D. B. Teplow, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 14745–14750.
- 38 N. G. Sgourakis, W.-M. Yau and W. Qiang, *Structure*, 2015, **23**, 216–227.
- 39 W. Qiang, W.-M. Yau, Y. Luo, M. P. Mattson and R. Tycko, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 4443–4448.
- 40 I. Bertini, L. Gonnelli, C. Luchinat, J. Mao and A. Nesi, *J. Am. Chem. Soc.*, 2011, **133**, 16013–16022.
- 41 K. Lindorff-Larsen, S. Piana, K. Palmo, P. Maragakis, J. L. Klepeis, R. O. Dror and D. E. Shaw, *Proteins: Struct., Funct., Bioinf.*, 2010, **78**, 1950–1958.
- 42 W. L. Jorgensen, J. Chandrasekhar, J. D. Madura, R. W. Impey and M. L. Klein, *J. Chem. Phys.*, 1983, **79**, 926–935.
- 43 S. T. Ngo, X.-C. Luu, M. T. Nguyen, C. N. Le and V. V. Vu, *RSC Adv.*, 2017, **7**, 42379–42386.
- 44 S. T. Ngo, H. M. Hung, N. D. Hong and N. T. Tung, *J. Mol. Graphics Modell.*, 2018, **83**, 122–128.
- 45 M. J. Abraham, T. Murtola, R. Schulz, S. Páll, J. C. Smith, B. Hess and E. Lindahl, *SoftwareX*, 2015, **1–2**, 19–25.
- 46 T. Darden, D. York and L. Pedersen, *J. Chem. Phys.*, 1993, **98**, 10089–10092.
- 47 W. G. Touw, C. Baakman, J. Black, T. A. H. te Beek, E. Krieger, R. P. Joosten and G. Vriend, *Nucleic Acids Res.*, 2015, **43**, D364–D368.
- 48 E. G. Marklund, M. T. Degiacomi, C. V. Robinson, A. J. Baldwin and J. L. P. Benesch, *Structure*, 2015, **23**, 791–799.
- 49 J. Srinivasan, T. E. Cheatham, P. Cieplak, P. A. Kollman and D. A. Case, *J. Am. Chem. Soc.*, 1998, **120**, 9401–9409.
- 50 P. A. Kollman, I. Massova, C. Reyes, B. Kuhn, S. Huo, L. Chong, M. Lee, T. Lee, Y. Duan, W. Wang, O. Donini, P. Cieplak, J. Srinivasan, D. A. Case and I. T. E. Cheatham, *Acc. Chem. Res.*, 2000, **33**, 889–897.
- 51 H. K. Srivastava and G. N. Sastry, *J. Chem. Inf. Model.*, 2012, **52**, 3088–3098.
- 52 O. Trott and A. J. Olson, *J. Comput. Chem.*, 2010, **31**, 455–461.
- 53 S. T. Ngo and M. S. Li, *Mol. Simul.*, 2013, **39**, 279–291.
- 54 G. M. Morris, R. Huey, W. Lindstrom, M. F. Sanner, R. K. Belew, D. S. Goodsell and A. J. Olson, *J. Comput. Chem.*, 2009, **30**, 2785–2791.
- 55 V. Knecht, *J. Phys. Chem. B*, 2010, **114**, 12701–12707.
- 56 S. T. Ngo, H. M. Hung, K. N. Tran and M. T. Nguyen, *RSC Adv.*, 2017, **7**, 7346–7357.
- 57 J. A. Lemkul, J. Huang and A. D. MacKerell, *J. Phys. Chem. B*, 2015, **119**, 15574–15582.
- 58 C. Haupt, J. Leppert, R. Röncke, J. Meinhardt, J. K. Yadav, R. Ramachandran, O. Ohlenschläger, K. G. Reymann, M. Görlach and M. Fändrich, *Angew. Chem., Int. Ed.*, 2012, **51**, 1576–1579.
- 59 M. D. Kirkitadze, M. M. Condrón and D. B. Teplow, *J. Mol. Biol.*, 2001, **312**, 1103–1119.
- 60 Y. Fezoui and D. B. Teplow, *J. Biol. Chem.*, 2002, **277**, 36948–36954.
- 61 L. Tran, N. Basdevant, C. Prévost and T. Ha-Duong, *Sci. Rep.*, 2016, **6**, 21429.
- 62 L. Tran, S. T. Ngo and M. T. Nguyen, *Chem. Phys. Lett.*, 2018, **696**, 55–60.
- 63 S. T. Ngo, X.-C. Luu, N. T. Nguyen, V. V. Vu and H. T. T. Phung, *PLoS One*, 2018, **13**, e0204026.
- 64 A. Baumketner, M. G. Krone and J.-E. Shea, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 6027–6032.
- 65 E. D. Agdeppa, V. Kepe, A. Petric, N. Satyamurthy, J. Liu, S. C. Huang, G. W. Small, G. M. Coled and J. R. Barrio, *Neuroscience*, 2003, **117**, 723–730.
- 66 C. A. Mathis, Y. M. Wang, D. P. Holt, G. F. Huang, M. L. Debnath and W. E. Klunk, *J. Med. Chem.*, 2003, **46**, 2740–2754.
- 67 D. Alagille, H. DaCosta, R. M. Baldwin and G. D. Tamagnan, *Bioorg. Med. Chem. Lett.*, 2011, **21**, 2966–2968.
- 68 Z. P. Zhuang, M. P. Kung, A. Wilson, C. W. Lee, K. Plossl, C. Hou, D. M. Holtzman and H. F. Kung, *J. Med. Chem.*, 2003, **46**, 237–243.
- 69 C.-W. Lee, Z.-P. Zhuang, M.-P. Kung, K. Plössl, D. Skovronsky, T. Gur, C. Hou, J. Q. Trojanowski, V. M. Y. Lee and H. F. Kung, *J. Med. Chem.*, 2001, **44**, 2270–2275.
- 70 Z. P. Zhuang, M. P. Kung, C. Hou, D. M. Skovronsky, T. L. Gur, K. Plossl, J. Q. Trojanowski, V. M. Y. Lee and H. F. Kung, *J. Med. Chem.*, 2001, **44**, 1905–1914.
- 71 E. K. Ryu, Y. S. Choe, K. H. Lee, Y. Choi and B. T. Kim, *J. Med. Chem.*, 2006, **49**, 6111–6119.
- 72 S. T. Ngo and M. S. Li, *J. Phys. Chem. B*, 2012, **116**, 10165–10175.
- 73 P. D. Huy, Y. C. Yu, S. T. Ngo, T. V. Thao, C. P. Chen, M. S. Li and Y. C. Chen, *Biochim. Biophys. Acta, Gen. Subj.*, 2013, **1830**, 2960–2969.
- 74 R. A. Rodriguez, L. Y. Chen, G. Plascencia-Villa and G. Perry, *ACS Chem. Neurosci.*, 2018, **9**, 783–789.

