

Small-Angle X-ray Scattering Studies of the Interaction of Amyloid Peptide Monomers with DPPC Oligo-Lamellar Vesicles

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The β -amyloid peptide ($A\beta$) is a 39-43 amino acid peptide that is a fragment derived from proteolytic cleavage of the large amyloid precursor protein (APP). The monomer of $A\beta$ is soluble, but at higher concentration $A\beta$ will self-assemble into fibril. The fibril aggregates of $A\beta$ are associated to Alzheimer's disease. The interaction between $A\beta$ and membrane is the key factor to the Alzheimer's disease. When $A\beta$ transforms into to β -sheet structure, $A\beta$ will aggregate to form fibrils and deposit on membrane. We have shown previously that both the $A\beta$ monomer and fibrils can interact with the multilamellar DPPC liposomes. In the current study, the interaction between the $A\beta$ monomer and the DPPC oligo-lamellar vesicles was investigated by SAXS.

Figure 1 shown the SAXS profiles of pure DPPC vesicles at different temperatures. The correlation peaks at $0.1 \text{ \AA}^{-1}$ became a broad peak as the temperature was increased to $43 \text{ }^\circ\text{C}$. At $43 \text{ }^\circ\text{C}$, the hydrocarbon tails were in liquid crystalline state (melted) and the correlation between the membranes within the multilamellar vesicles were destroyed. Only the form factor of the bilayer was observed at $43 \text{ }^\circ\text{C}$.

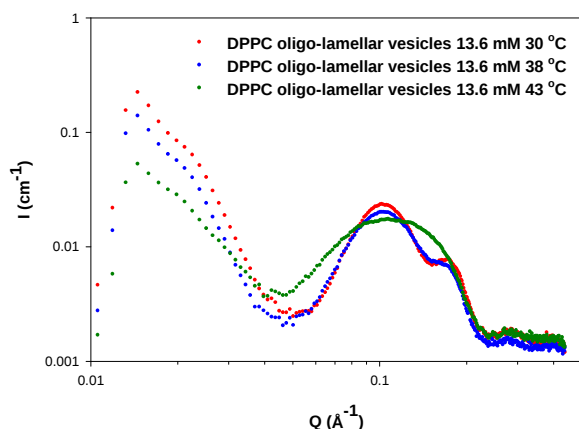


Figure 1. The SAXS profiles of DPPC oligo-lamellar vesicles at $30 \text{ }^\circ\text{C}$, $38 \text{ }^\circ\text{C}$ and $43 \text{ }^\circ\text{C}$.

Figure 2 shows the effect of adding $A\beta$ monomers at $30 \text{ }^\circ\text{C}$. The scattering profiles are similar but the scattering intensity is higher when more $A\beta$ monomers are added. It is evident that $A\beta$ monomers are indeed inserted into the DPPC bilayers.

The scattering profiles of DPPC oligo-lamellar vesicles inserted with $A\beta$ monomers at $43 \text{ }^\circ\text{C}$ are shown

in Figures 3. Similar to the pure DPPC case, at $43 \text{ }^\circ\text{C}$ only the form factor of the bilayer can be observed. With the insertion of $A\beta$, the scattering intensity is higher due to the incorporation of the $A\beta$ in the DPPC bilayer. Detail analyses will be carried out to reveal the detail structure of the DPPC bilayers inserted with $A\beta$.

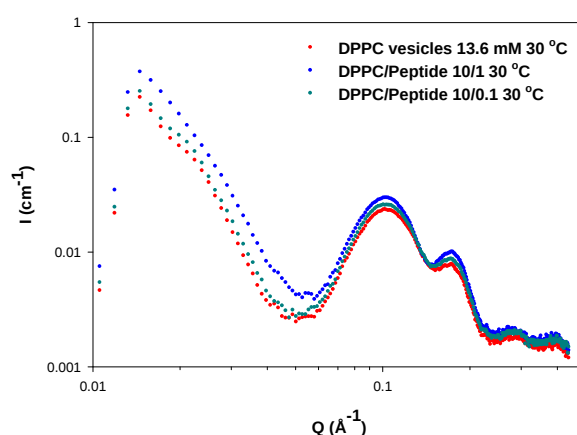


Figure 2. DPPC oligo-lamellar vesicles with $A\beta$ monomers at $30 \text{ }^\circ\text{C}$. The lipid to peptide ratios were equal to 10/1 and 10/0.1 respectively.

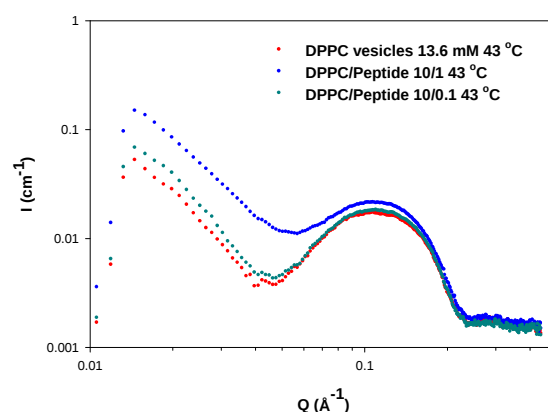


Figure 3. DPPC oligo-lamellar vesicles with $A\beta$ monomers at $43 \text{ }^\circ\text{C}$. The lipid to peptide ratios were equal to 10/1 and 10/0.1 respectively.