

Supporting information

Ruthenium complexes bearing glucosyl ligands are able to inhibit the amyloid aggregation of short Histidine-peptides

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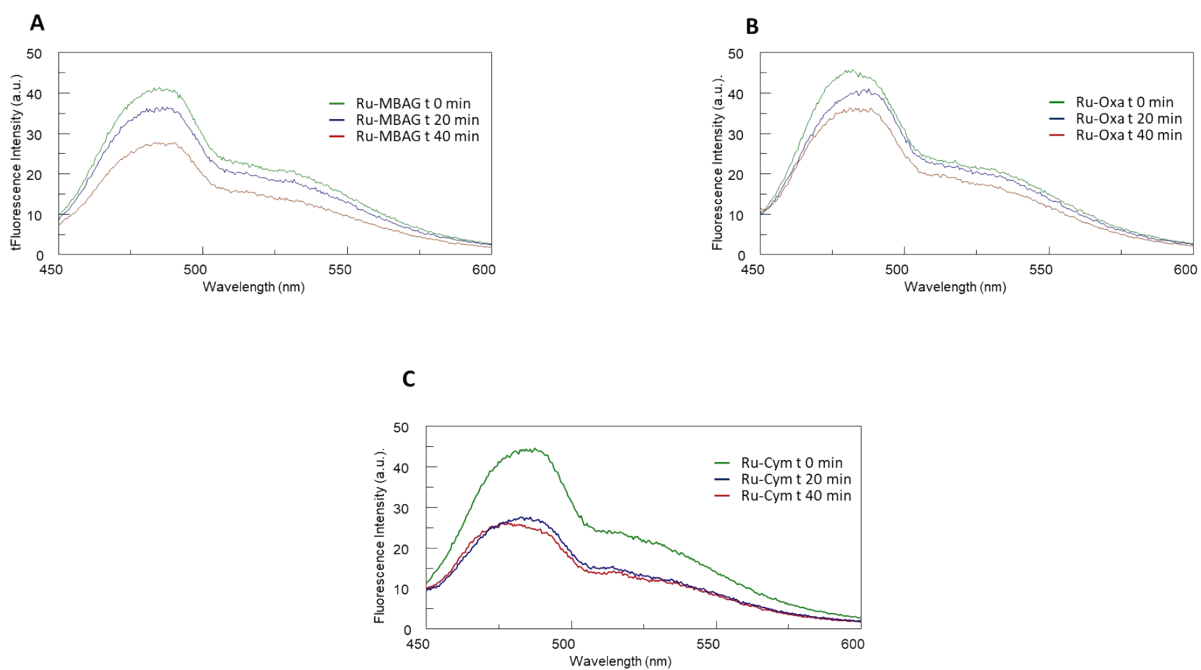


Fig. S1 Overlay of ThT fluorescence emission intensity of Ru-MBAG (A), Ru-Oxa (B) and Ru-Cym (C) complexes at indicated time.

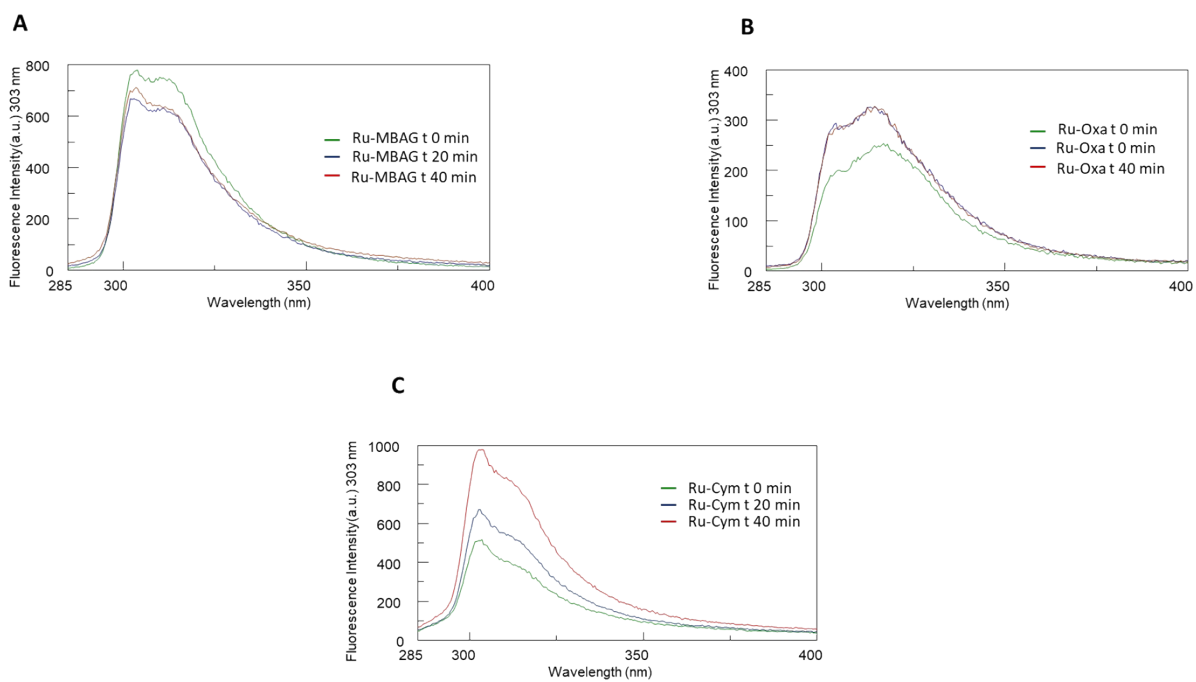


Fig. S2 Overlay of aromatic intrinsic fluorescence assay of Ru-MBAG (A), Ru-Oxa (B) and Ru-Cym complexes (C) at indicated time.

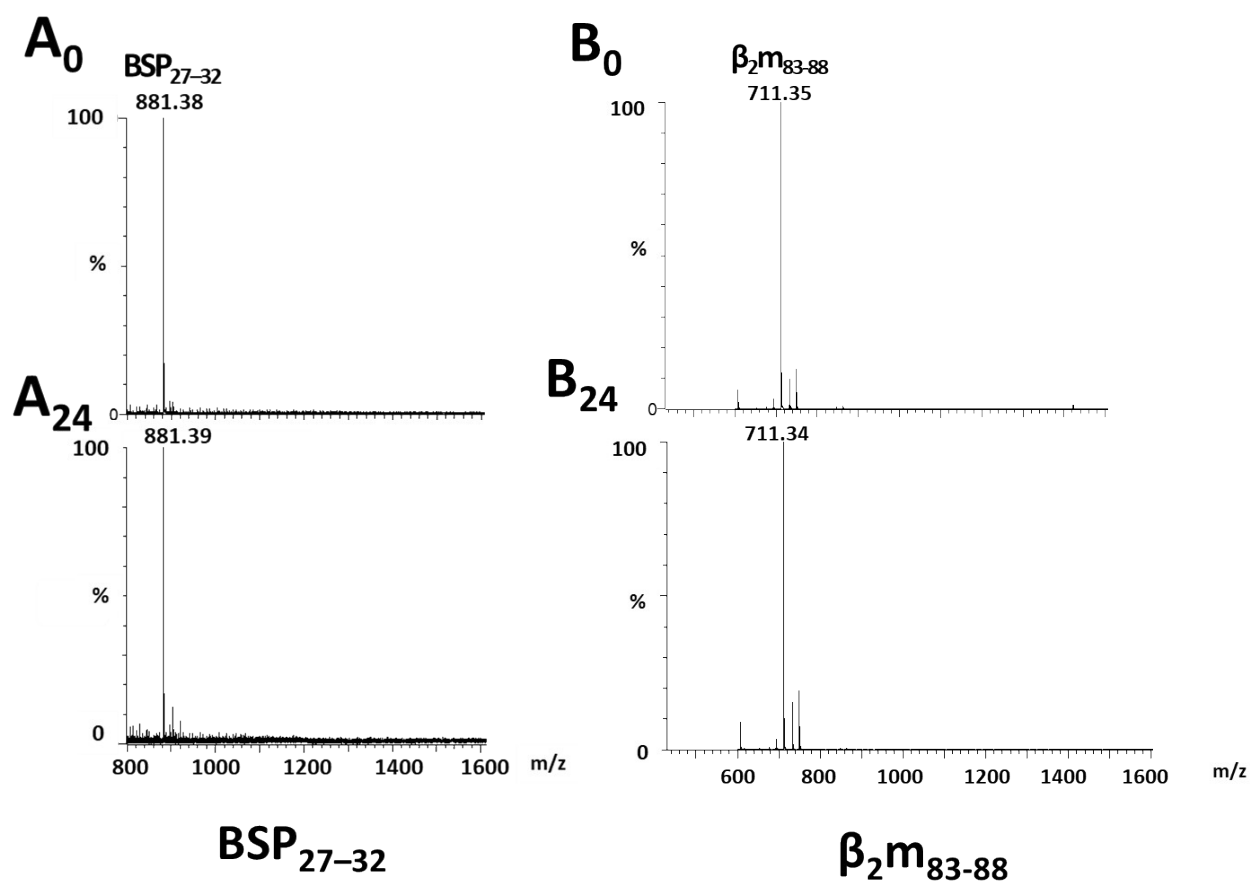


Figure S3. ESI-MS spectra of isolated BSP_{27-32} (A₀, A₂₄ panels) and β_2m_{83-88} (B₀, B₂₄ panels) at 0 and 24h

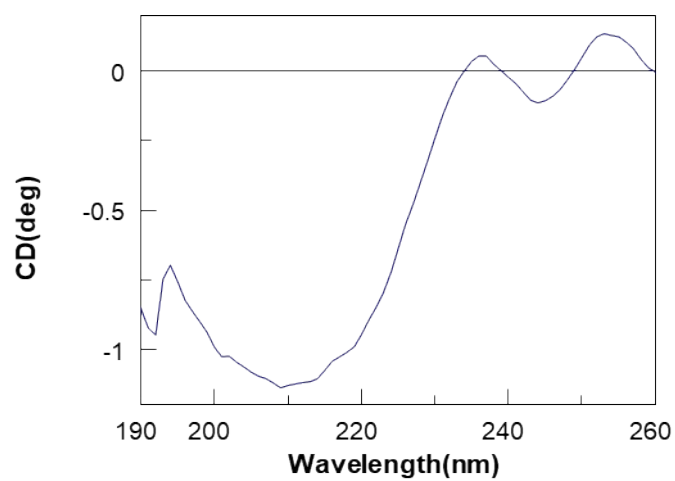


Figure S4. Conformational analysis of BSP_{27-37} at 1mM, phosphate buffer 10mM.

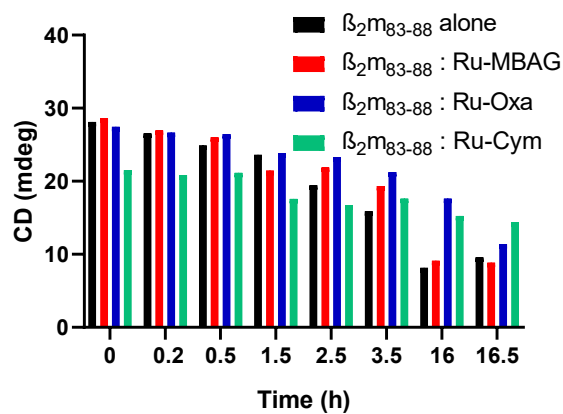


Figure S5. Histogram of CD absolute values of β_2m_{83-88} at indicated times.

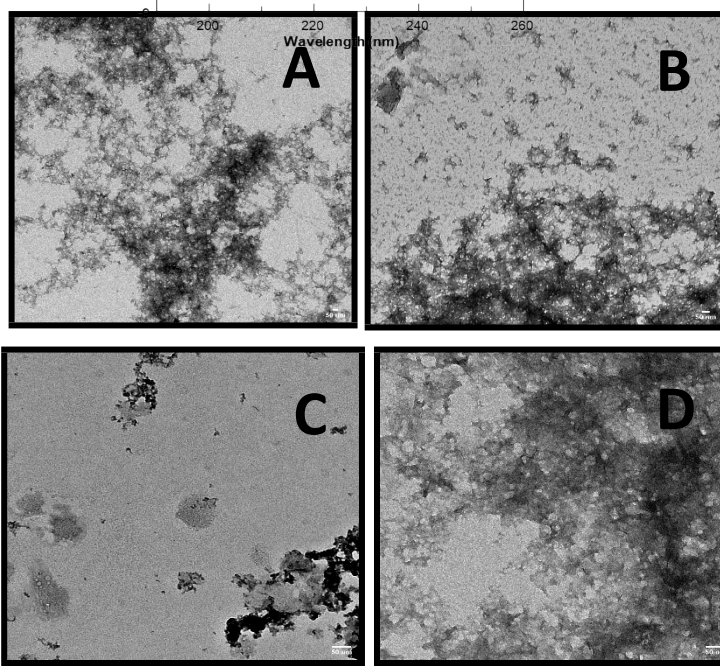
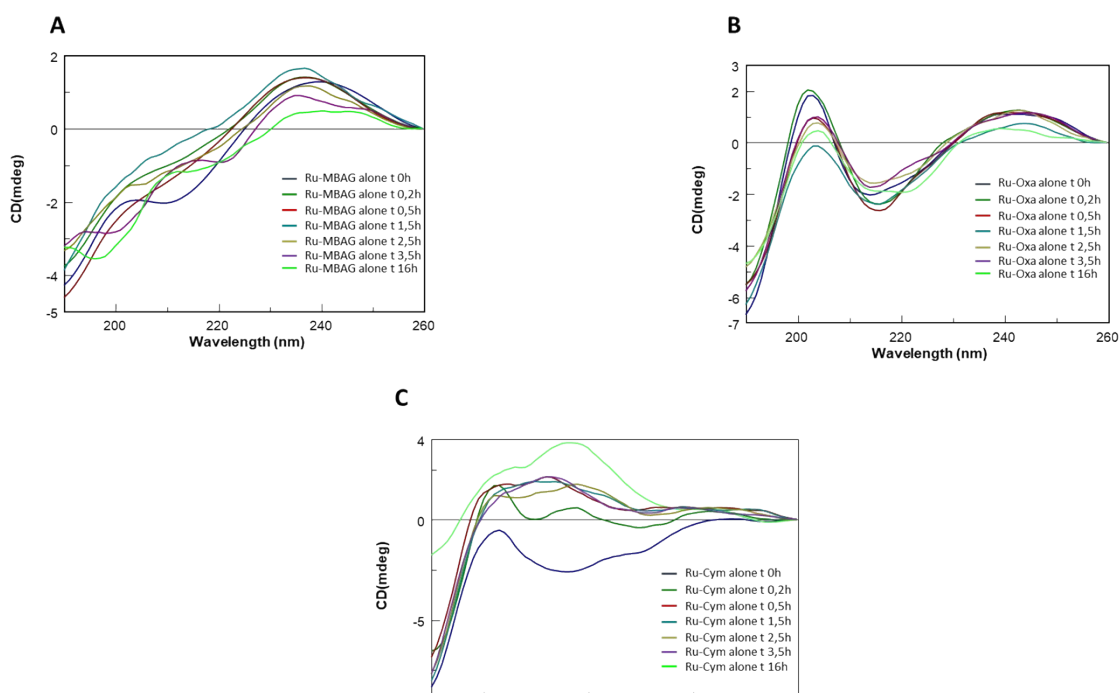


Fig. S6
analysis of Ru(II)-
Ru-MBAG alone
alone (400 μ M) and
(400 μ M).

Conformational
arene complexes. (A)
(400 μ M), (B) Ru-Oxa
(C) Ru-Cym alone

Figure S7. TEM analysis at t=4 d of BSP₂₇₋₃₂ peptide (A) alone, and in presence of (B) Ru-Cym, (C) Ru-MBAG and (D) Ru-Oxa.

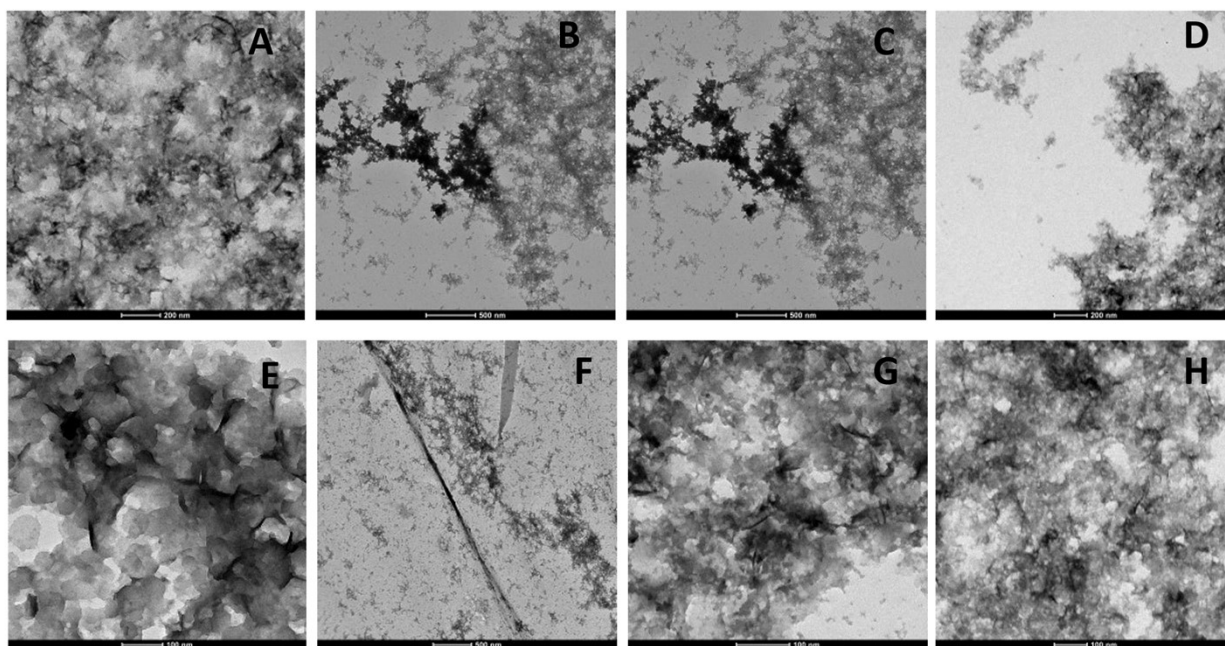


Fig. S8 TEM analysis of $\beta 2m_{83-88}$ peptide alone at t 0 and 4 d (A and E) and in presence of Ru-Cym, Ru-MBAG and Ru-Oxa (B-D, t=0), (F-H, t= 4 d).