Supporting Information

Controlled and Selective Photo-oxidation of Amyloid-β Fibrils by Oligomeric *p*-Phenylene Ethynylenes

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Figure S1: Reverse phase HPLC chromatographs of soluble A β 40 before and after irradiation in the presence of MB or OPE₁²⁻. Unincubated A β 40 monomers, whether non-irradiated or irradiated in the presence of OPE₁²⁻, display a similar elution time that corresponds to 44.6-44.4% acetonitrile. When A β 40 monomers are irradiated in the presence of MB, the elution profile of A β 40 monomers significantly changed. The main peak eluted earlier (43.5% acetonitrile), which shows that A β 40 peptide became more hydrophilic, which is consistently with the oxidation of the peptide. Also, the elution profile is broad with multiple peaks, indicating the presence of a several populations of A β 40. Experimental method: The monomeric protein was analyzed by RP-HPLC on an Agilent 1100 instrument (Agilent Technology, Santa Clara, CA) before and after irradiation in the presence of OPE₁²⁻ or MB (5 μ M protein with 1 μ M photosensitizer). 110 μ L of 5 μ M protein was centrifuged at 14,000 rpm for 15 minutes. The supernatant (100 μ L) was loaded onto an Eclipse XDB C18 column (Agilent Technology, Santa Clara, CA) pre-equilibrated at 40 °C with 95% of mobile phase A (water containing 0.1% TFA) and 5% of mobile phase B (acetonitrile containing 0.1% TFA). A β 40 was eluted using a 5-100% linear gradient of mobile phase B over

40 min. The absorbance at 215 nm was monitored. Each chromatogram was background subtracted using the Agilent ChemStation software.



Figure S2: TEM images of A β 40 after incubation of 50 μ M monomers either alone (A) or in the presence of 2.6 μ M non-sonicated A β 40 fibrils (B) or 2.6 μ M sonicated A β 40 fibrils (C). Only A β protofibrils produced by sonication promoted fast peptide fibrillation.