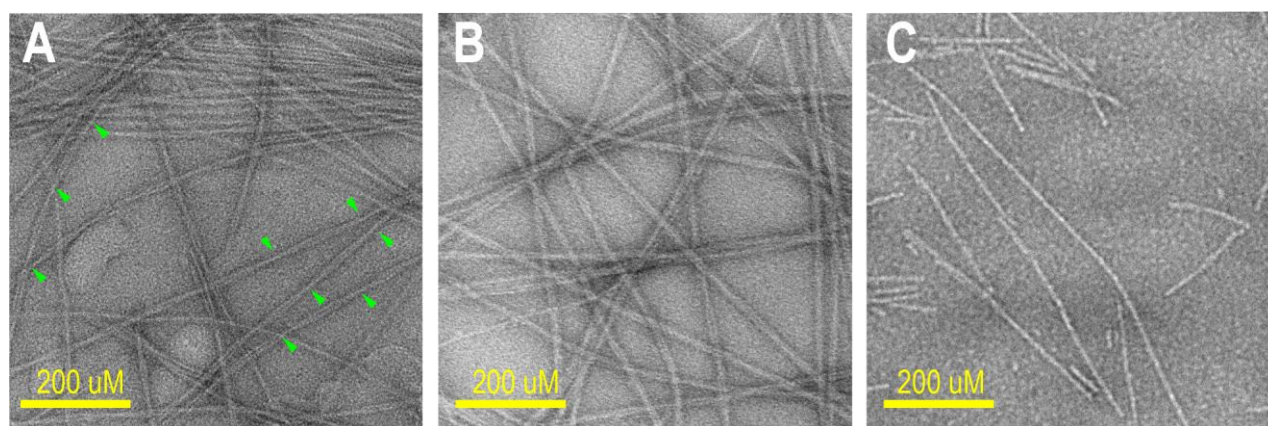
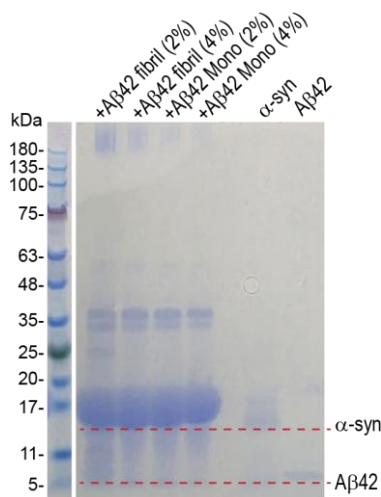


**Fig. 1.** (A) SDS page gel image and (B) size exclusion chromatography of the purified  $\alpha$ -syn monomer. (C) ThT fluorescence intensities of  $\alpha$ -syn alone (black), with A $\beta$ 42 fibrils (red), or A $\beta$ 42 monomers (orange) at an  $\alpha$ -syn: A $\beta$ 42 ratio of 50:1.



**Fig. 2.** Negatively stained TEM images of  $\alpha$ -syn fibrils formed in the presence of A $\beta$ 42 fibrils (A), A $\beta$ 42 monomers (B), or in the absence of A $\beta$ 42 (C). Green arrows in (A) indicate twisted fibrils.



**Fig. 3.** SDS page analysis of protein aggregates. From left to right:  $\alpha$ -syn fibrils formed with A $\beta$ 42 fibril and A $\beta$ 42 monomers (2 and 4 % relative to  $\alpha$ -syn),  $\alpha$ -syn fibrils alone, and A $\beta$ 42 fibrils alone. Distinct bands corresponding to  $\alpha$ -syn ( $\sim$ 14kDa) and A $\beta$ 42 ( $\sim$ 5kDa) are indicated with red dashed lines, confirming the presence of both proteins in the mixed aggregates.