Supplementary data

Amyloid–β 1–42 aggregation initiates its cellular uptake and cytotoxicity

Sha Jin^{1, 2}, Niraja Kedia²,Eva Illes—Toth², Ivan Haralampiev³, Simon Prisner³, Andreas Herrmann⁴, Erich E. Wanker¹ and Jan Bieschke²

Supplementary Figure 1: Live cell FRET imaging of $A\beta$ internalization. SHEP cells were incubated with fibrillar $A\beta_{1-42}^{633}$ (150 nM) for 30 min followed by monomeric $A\beta_{1-42}^{555}$ (150 nM). Co-aggregation of monomeric and fibrillar $A\beta_{1-42}$ was quantified by FRET analysis. Beta-sheet structures were co-stained with ThS. Shown is one representative time course out of four independent experiments.

¹Proteomics and Molecular Mechanisms of Neurodegenerative Diseases, Max Delbrück Center for Molecular Medicine, Berlin–Buch, Germany

²Department of Biomedical Engineering, Washington University in St. Louis, MO, USA

³ Department of Biology, Humboldt–Universität zu Berlin, Berlin, Germany

⁴Department of Biology, IRI Life Sciences, Humboldt–Universität zu Berlin, Berlin, Germany

