Supporting Information for

Comparison of multiple reported Amyloid-ß receptors for sufficiency, affinity, and Alzheimer's relevance

Levi M. Smith^{1,2}, Mikhail A. Kostylev¹, Suho Lee¹, and Stephen M. Strittmatter^{1,*}

From the ¹Program in Cellular Neuroscience, Neurodegeneration & Repair, Departments of Neurology and of Neuroscience, Yale University School of Medicine, New Haven, CT 06536, USA, and ²Department of Cell Biology, Yale University School of Medicine, New Haven, CT 06536, USA

*To whom correspondence should be addressed: Stephen M. Strittmatter, CNNR Program, BCMM 436, Yale University School of Medicine, 295 Congress Avenue, New Haven, CT 06536, USA stephen.strittmatter@yale.edu

This PDF includes:

Figure S1



Figure S1. Neuropilin-1 does not bind Aß, regardless of preparation.

A. COS-7 cells expressing the indicated Myc-tagged protein and incubated with 1 μ M BABo, BABm, or ABg at 4°C or BABo at 37°C. Scale bar = 200 μ m. BABo = biotinylated amyloid beta oligomers (o), monomers (m), or globulomers (g) **B**. Quantification of binding of different AB preparations to NRP1-expressing cells relative to BABo (or ABo for globulomer) binding to Myc-hPrP^C transfected cells. One-sided T test comparing to an expected value of 100 (% hPrP^C BABo or ABo binding at 4°C). N = 3 - 4 experiments.