## Blockade of dengue virus infection and viral cytotoxicity in neuronal cells *in vitro* and *in vivo* by targeting endocytic pathways

Min-Ru Ho, Tsung-Ting Tsai, Chia-Ling Chen, Ming-Kai Jhan, Cheng-Chieh Tsai, Yi-Chao Lee, Chun-Han Chen, and Chiou-Feng Lin

## Supplemental information

Supplemental Figure 1 DENV2 causes D2R-mediated infection. Flow cytometry and immunofluorescent analysis showed Neuro-2a (A) and primary cultured hippocampal neuronal cells (**B**) expressing D2R or D4R. (**C**) Primary cultured hippocampal neuronal cells were inoculated with Alexa-594 labeled (red) DENV2 (MOI=1) for 2 h. The fluorescent image shows cells carrying fluorescent DENV2. (D) Plaque assays showed the level of viral replication. Neuro-2a cells were infected with DENV2 (MOI=1) with or without D2R inhibitor MCP (10 µM) pre-treatment. (E) The fluorescent image shows cells carrying Alexa-594 labeled (red) DENV2 2 h post-infection. (F) Plaque assays show the level of viral replication. For all images and histograms, representative data were selectively obtained from three individual experiments. DAPI staining indicates the nuclei (*blue*). For flow cytometry analysis, the percentage of positive cells is shown. All quantitative data are shown as the means  $\pm$  SD from three independent experiments. \*p < 0.05, \*\*p < 0.01, and \*\*\*p < 0.001. ns, not significant.



## Ho et al Supplemental Figure 1