## **Supplementary Figure S9**



Supplementary Figure S9. TBS treatment rescues a synaptic loss in  $CA1_{PV}$  from AD mice. (a) Synaptophysin (SYT)-labeled presynaptic terminals (pink) in the CA1 hippocampus from control and AD mice, in which eGFP was expressed in the  $CA1_{PN}$  (green). (b, c) Presynaptic terminals (b) and postsynaptic spines (c) in the CA1<sub>PN</sub> were analyzed in the CA1 sl region from control and AD mice. The mice were treated with (AD/TBS) or without (AD/non-TBS). Data are mean  $\pm$  SEM (n = 4 mice per group, \*p < 0.001, t-tests). (d) Representative images of a CA1 hippocampus from control and AD mice that expressed eGFP in CA1<sub>PV</sub> cells. The mice were treated with (AD/TBS) or without (AD/non-TBS). Images show the spines in the PV dendritic branches in the CA1 sl region, a terminal projection zone of  $ECII_{PN}$ . (e) Spine densities in the CA1<sub>PV</sub> of AD are reduced and rescued by TBS treatment. Data are mean  $\pm$  SEM (n = 4 mice per group, \*p < 0.001, t-tests). In this study, AD/ECII<sub>PN</sub><sup>ChR2+</sup> (AD mice) mice were crossed with the CaMK-II $\alpha$ -Cre (a) or the PV-Cre (d) mice and stereotaxically injected with 2  $\mu$ l of a high titer of the lenti-eGFP<sup>loxP</sup>/loxP virus particles (3  $\times$  10<sup>11</sup> genomic particles/ml) in the CA1 region, resulting in eGFP expression in the CA1<sub>PN</sub> or the CA1<sub>PV</sub> of AD/ECII<sub>PN</sub><sup>ChR2+</sup> mice. Presynaptic terminals and the dendritic branches (20 µm segment) from the  $CA1_{PN}$  (50 segments per animal) and the  $CA1_{PV}$  (50 segments per animal) in the sl region were studied. The averaged spine densities (spines per 10  $\mu$ m<sup>2</sup> length of dendrites) was analyzed 25 days after 35-days TBS treatment.

so: stratum oriens; sp: stratum pyramidale; sr: stratum radiatum; sl: stratum lacumosum; sm: stratum moleculare.