

Figure S1. OEA treatment inhibits microglia M1 polarization in the peri-infarct of the cortex and striatum in WT but not KO mice at 3 days after MCAO. (A) Representative immunofluorescence images and quantification of TMEM119, iNOS and CD16/32 in the peri-infarction of WT mice on day 3 after MCAO. (B) Representative immunofluorescence images and quantification of TMEM119, iNOS and CD16/32 in the peri-infarction of KO mice on day 3 after MCAO. The data are the means \pm SEM. n = 5 per group. ** $p < 0.01$ vs MCAO + vehicle group; N.S = No Significance. Scale Bar=100 μ m.

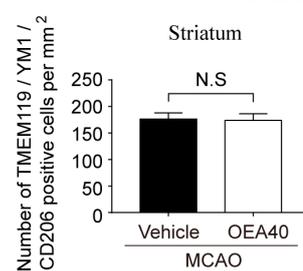
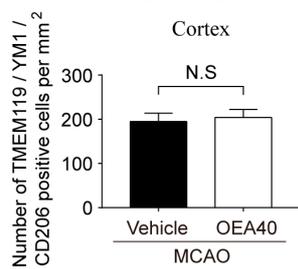
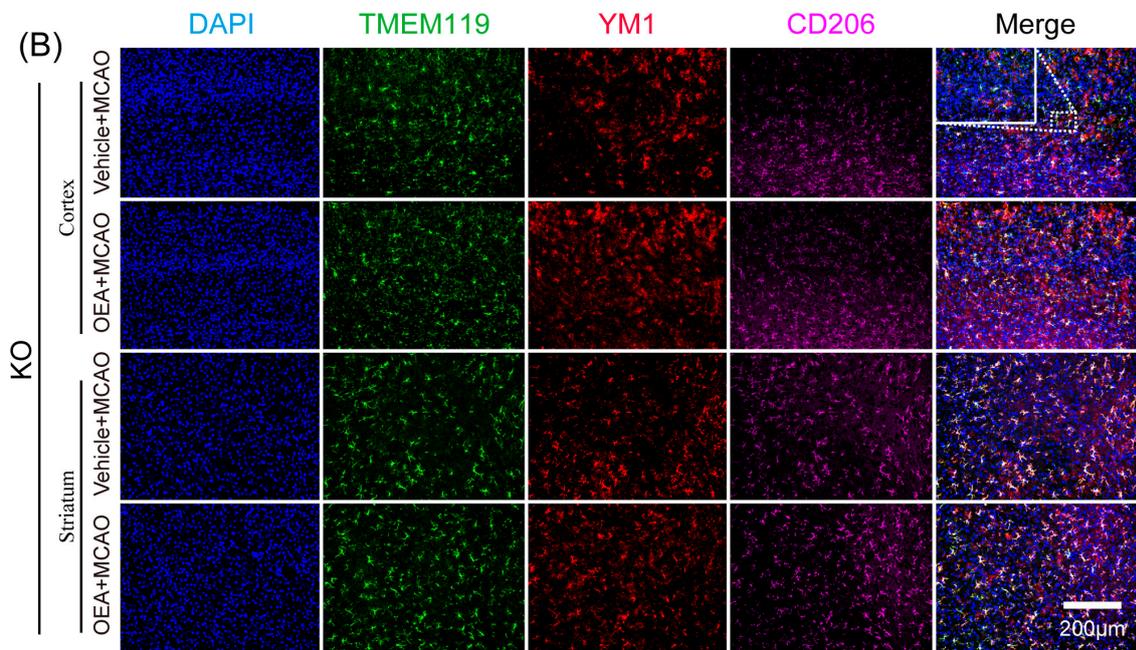
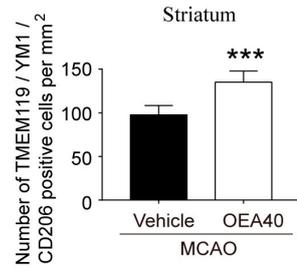
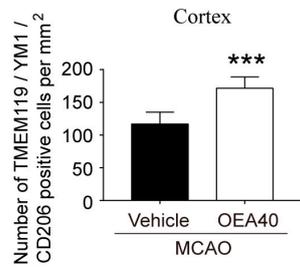
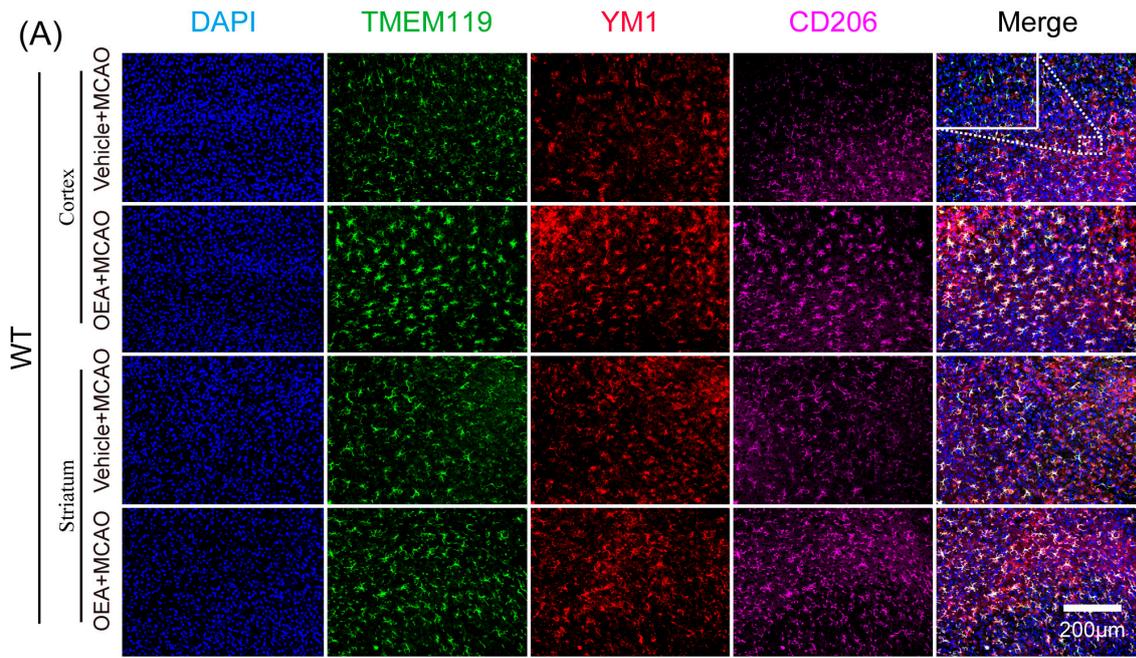


Figure S2. OEA treatment promotes microglia M2 polarization in the peri-infarct of the cortex and striatum in WT but not KO mice at 3 days after MCAO. (A) Representative immunofluorescence images and quantification of TMEM119, YM1 and CD206 in the peri-infarction of WT mice on day 3 after MCAO. (B) Representative immunofluorescence images and quantification of TMEM119, YM1 and CD206 in the peri-infarction of KO mice on day 3 after MCAO. The data are the means \pm SEM. $n = 5$ per group. $^{***}p < 0.001$ vs MCAO + vehicle group; N.S = No Significance. Scale Bar=100 μ m.

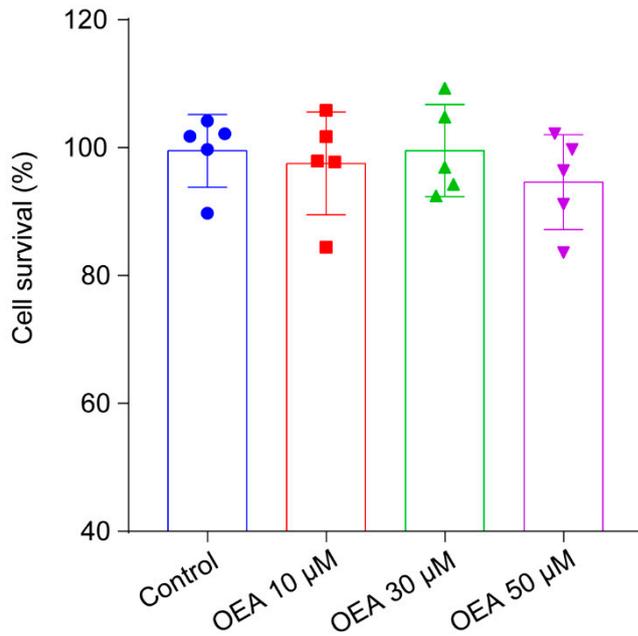


Figure S3. Effects of OEA on BV2 cells viability. BV2 cells were cultured with different concentrations of OEA (10, 30, and 50 μ M) for 24 h. Cell viability was assessed using an MTT assay. The values are presented as means \pm SEM of five independent experiments performed in duplicate ($n = 5$). $^{*}p < 0.05$ vs. control group.

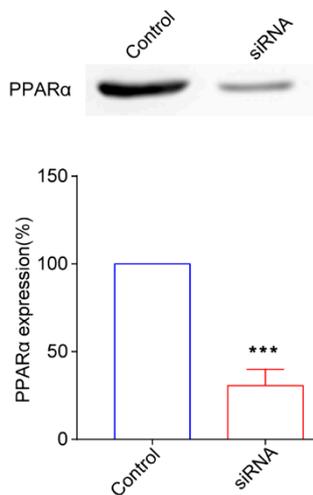


Figure S4. BV2 cells were transiently transfected with PPAR α siRNA for 24 h, and the protein expression of PPAR α were subsequently measured by western blot. The values are presented as means \pm SEM of five independent experiments performed in duplicate ($n = 5$). $^{***}p < 0.001$ vs. control group.

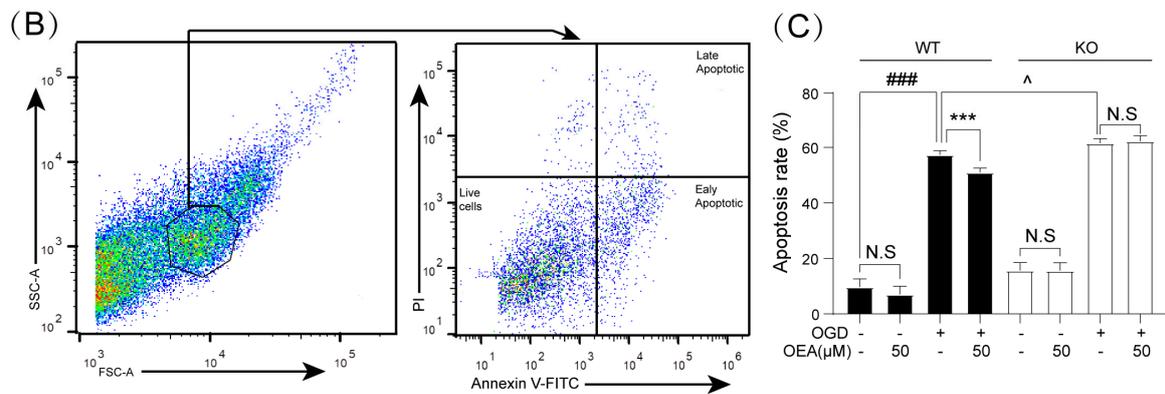
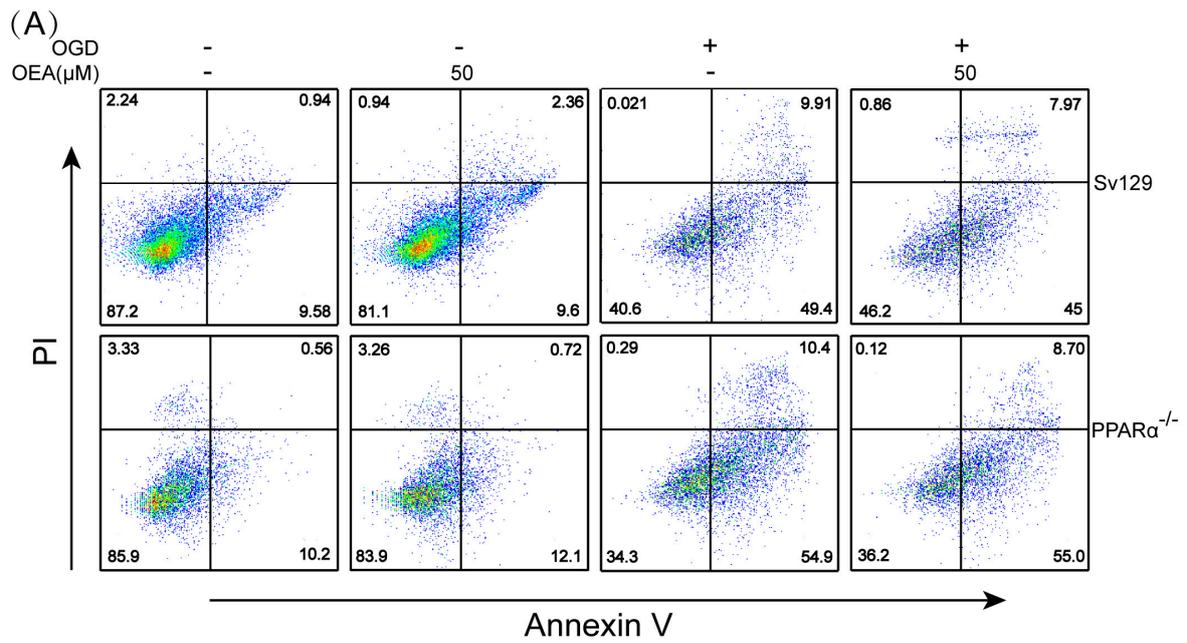


Figure S5. OEA treatment protects against neuronal apoptosis after OGD-treated in N/G co-cultures through PPAR α . (A) Neuronal apoptosis was assessed with flow cytometry using annexin V-FITC staining. (B and C) The relative apoptosis ratio was analyzed. The data are presented as the mean \pm SEM. N = 5. Each experiment was repeated 3 times. ### p < 0.001 vs. Control group, *** p < 0.001 vs. OGD group, ^ p < 0.05 vs. WT OGD group