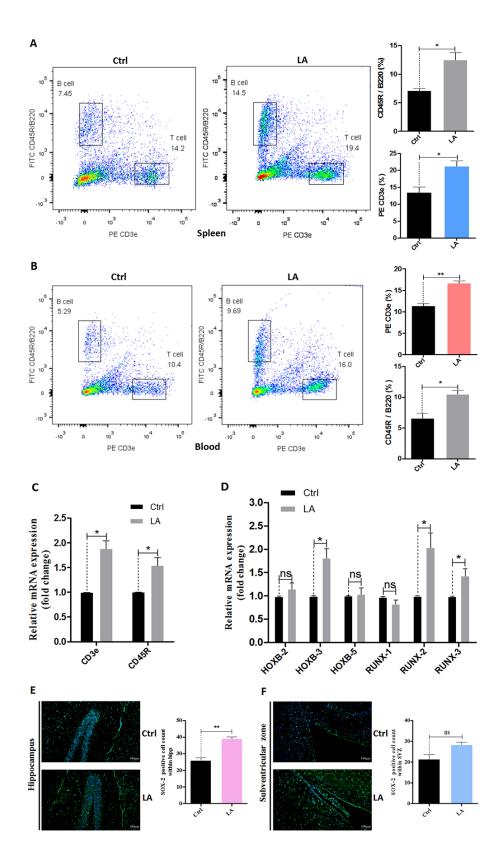
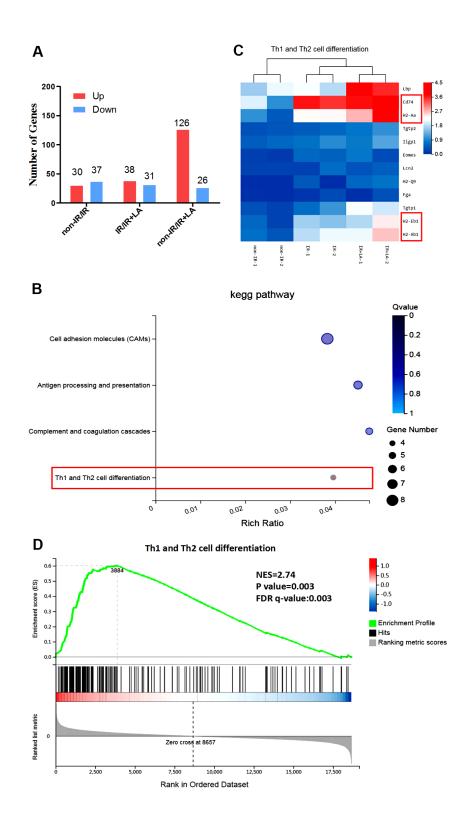


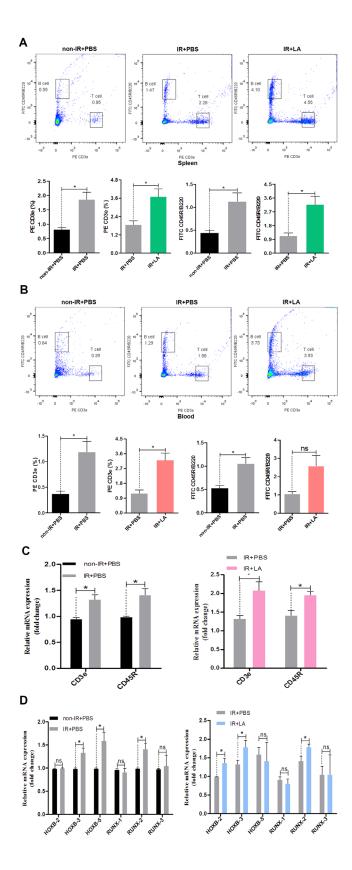
Supplementary Figure 1. LA effect on effector T-and B-cell proliferation in the spleen and whole blood in C57BL/6 mice. (A) Effector T-cell proliferation in the spleen of LA-or control (Ctrl)-treated mice. (B) Effector B-cell proliferation in the spleen of LA-or control (Ctrl)-treated mice. (C) Effector T-cell proliferation in the whole blood of LA-or control (Ctrl)-treated mice. (D) Effector B-cell proliferation in the whole blood of LA-or control (Ctrl)-treated mice. The data are presented as the mean \pm SD of three independent experiments. Statistical significance was determined using unpaired t-tests. *p < 0.05, **p < 0.01 compared with the Ctrl group.



Supplementary Figure 2. Effect of LA on lymphocyte proliferation in vitro. (A, B) Proliferation of LA-treated or control (Ctrl)-treated in vitro cultured T and B cells from the spleen (A) or whole blood (B) of C57BL/6 mice. (C) Relative mRNA expression of in vitro splenic lymphocytes. (D) Relative mRNA expression of genes that might regulate lymphocyte proliferation. (E, F) Expression of the neural stem cell marker SOX-2 in the hippocampus (Hipp) (E) and SVZ (F) of C57BL/6 mice. The data are presented as the mean±SD of three independent experiments. Statistical significance was determined using unpaired t-tests. *p < 0.05, **p < 0.01, as compared with the Ctrl.



Supplementary Figure 3. Exploring the mechanism by which LA regulates the immune system by performing RNA-seq on the hippocampus of B-NDG mice. (A) The differentially expressed genes between non-IR and IR PBS-treated mice, between non-IR PBS-treated mice and IR LA-treated (IR+LA) mice, and between IR PBS-treated mice and IR+LA mice. (B) A bubble diagram showing the Th1- and Th2-cell differentiation signaling pathway regulated by LA treatment. (C) Heatmap showing the differentially expressed genes related to the Th1- and Th2-cell differentiation signaling pathway. (D) Enrichment plots of gene expression signatures for Th1- and Th2-cell differentiation. Barcode plot indicates the positions of genes in each gene-set. NES, normalized enrichment score. The data are presented as the mean \pm SD of three independent experiments. Statistical significance was determined using unpaired t-tests.*p < 0.05, **p < 0.01, as compared with the control.



Supplementary Figure 4. The effect of LA on the *in vitro* proliferation of lymphocytes from B-NDG mice. (A, B) Proliferation of LA-treated or PBS-treated *in vitro* cultured T and B cells from the spleen (A) or whole blood (B) of IR B-NDG mice. (C) The relative mRNA expression of CD3e and CD45R. (D) Relative mRNA expression of genes that might regulate lymphocyte proliferation. The data are presented as the mean±SD of three independent experiments. Statistical significance was determined using unpaired t-tests.*p < 0.05, **p < 0.01, as compared with the Ctrl.