SUPPLEMENTARY FIGURE LEGENDS

Supplementary Figure 1. Detection of SREBP2 mature form and precursor form in HepG2 cells without and with ANA treatment

HepG2 cells were treated with vehicle DMSO or 5 μ M ANA for 24 h. Total cell lysates were isolated and protein levels of SREBP2 were examined by Western blotting. SREBP2-P refers to the precursor form and SREBP-M refers to the processed and active mature form of SREBP2.

Supplementary Figure 2. Mice serum lipid levels

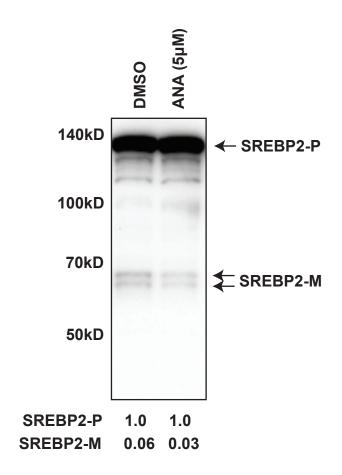
Male C57BL/6J mice (n=16) were fed a normal diet for 1 week and then switched to a HFHC diet for two weeks. Four hour-fasted serum lipids were measured before and after the two week HFHC feeding.

Supplementary Figure 3. ANA treatment affected serum cholesterol levels and LDLR/PCSK9 expression in mice fed a normal chow diet.

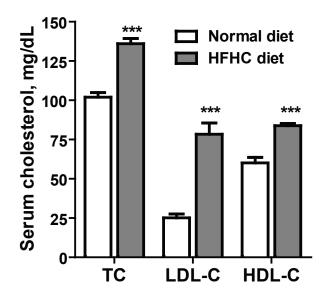
- (A) Male C57BL/6J mice fed a normal chow diet were orally dosed 50 mg/kg/day ANA (n=8) or equal volume of vehicle (0.5% methyl cellulose) as the control group (n=8) for 10 days. Serum TC, HDL-C and LDL-C were measured at day 10 after the last dosing by commercial kits. Data are mean ±SEM. Significant differences between control and ANA treatment were assessed by two-tailed Student's t-test.
- (**B**) qPCR analysis of liver mRNA levels of LDLR and PCSK9 along with 4 additional SREBP-target genes and 3 LXR regulated genes in ANA-treated and vehicle-treated control mice. Values are mean \pm SEM of 8 mice per group. *p < 0.05 and **p < 0.01 compared to the vehicle group.

- (C) Individual liver protein extracts were prepared and protein concentrations were determined. 50 μg of homogenate proteins of individual liver samples were resolved by SDS-PAGE and LDLR protein was detected by immunoblotting using anti-LDLR antibody. The membrane was reprobed with anti-β-actin antibody. Individual nuclear extracts isolated from liver homogenate from the ANA and vehicle groups were analyzed for SREBP2-M protein levels by Western blotting. The membrane was reprobed with anti-HDAC1 antibody as a control of equal nuclear protein loading.
- (**D**) The protein abundance of LDLR was quantified with the Alpha View Software with normalization by signals of β -actin. Values are mean \pm SEM of 8 samples per group. The protein abundance of SREBP2-M was quantified with normalization by signals of HDAC1. Values are mean \pm SEM of 8 samples per group.
- (E) Individual serum PCSK9 levels were quantified by ELISA. Values are mean \pm SEM of 8 mice per group. * p < 0.05 as compared to vehicle group.
- (F) Cholesterol levels in vehicle and ANA-treated liver samples were measured. Values are mean \pm SEM of 8 mice per group. ** p < 0.01 as compared to vehicle group.

Supplementary Figure 1



Supplementary Figure 2



Supplementary Figure 3

