

Figure S1

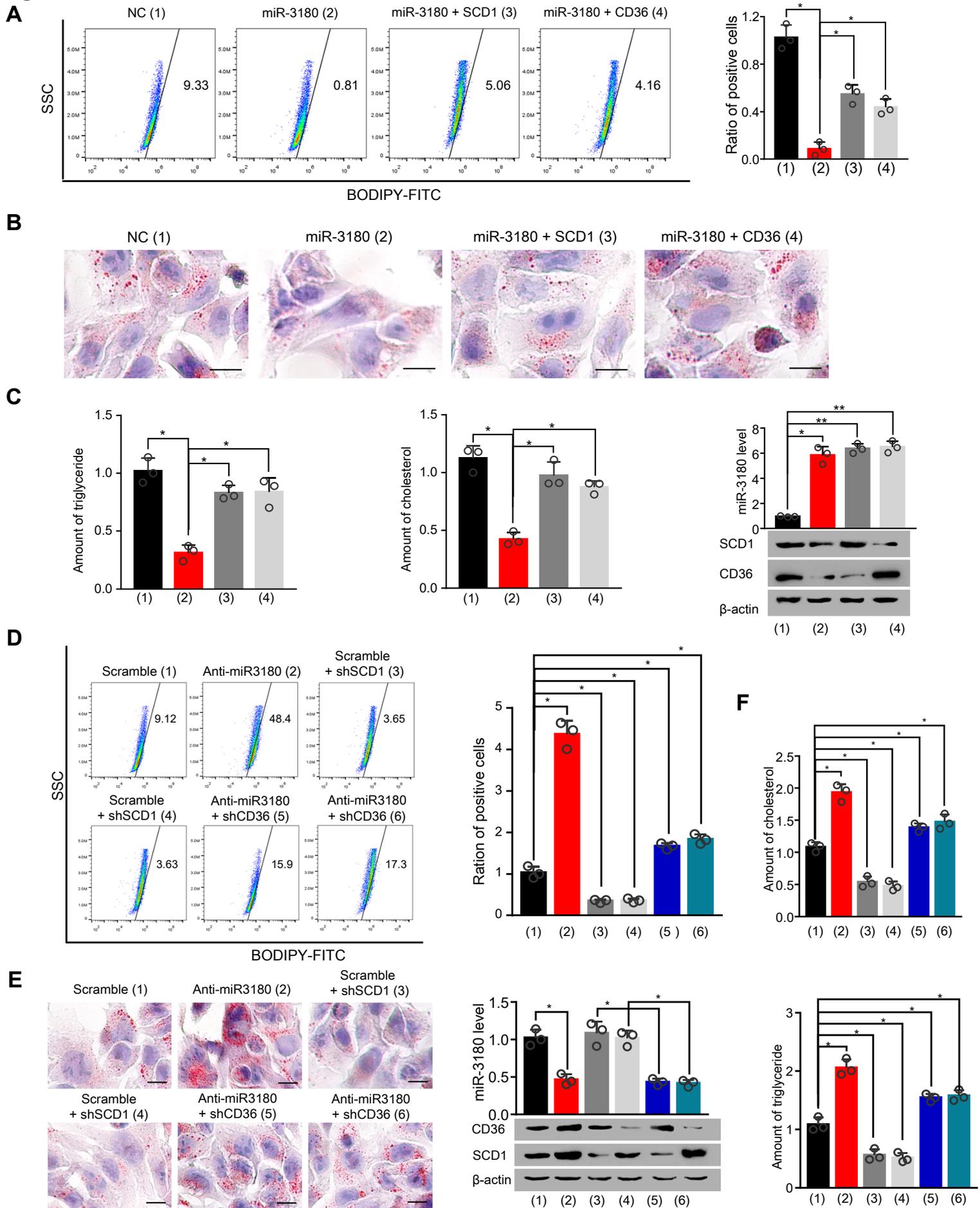
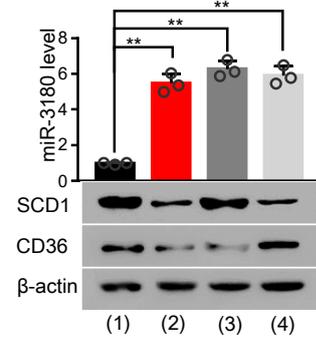
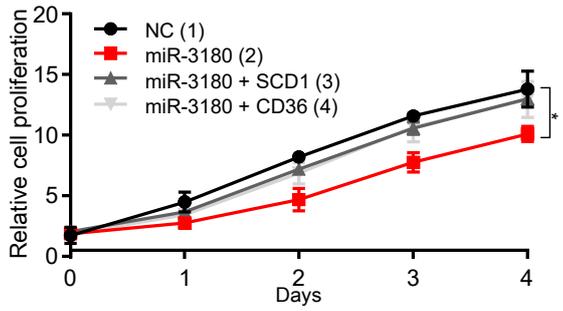


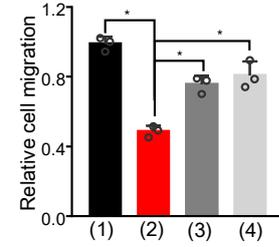
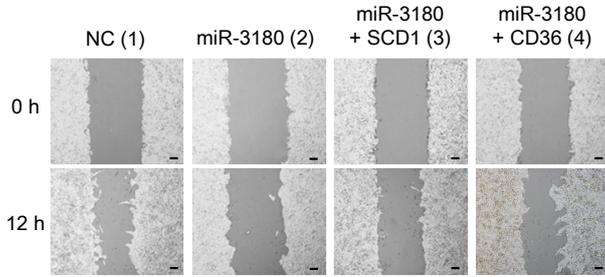
Figure S1. MiR-3180 suppresses lipid content in CD36- and SCD1-dependent manner. (A) Flow cytometry analysis of lipid content in MHCC-97H cells transfected with miR-3180 and the indicated expression vectors or negative control. The relative of lipid content are shown in the right panel. (B) Oil red O staining of lipid content in MHCC-97H cells transfected as (A). Scale bar: 20 μ m. (C) The relative triglyceride, cholesterol and the expression of miR-3180, SCD1 and CD36 levels in MHCC-97H cells transfected as (A). (D) Flow cytometry analysis of lipid content in MHCC-97H cells transfected with the indicated vectors. The relative lipid content is shown in the right nearby panel. (E) Oil red O staining of lipid content, RT-qPCR and immunoblotting analysis of miR-3180, CD36 and SCD1 expression in MHCC-97H cells transfected as (D). Scale bar: 20 μ m. (F) Relative triglyceride and cholesterol level in MHCC-97H cells transfected as (D). Data shown are presented as mean \pm SD of triplicate measurements with similar results. **P < 0.01.

Figure S2

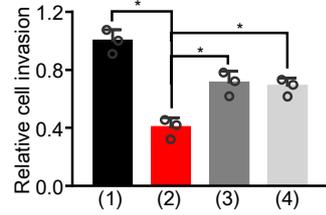
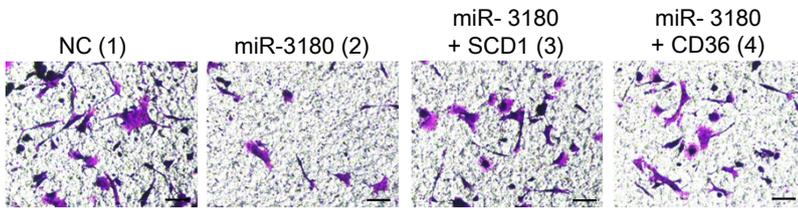
A



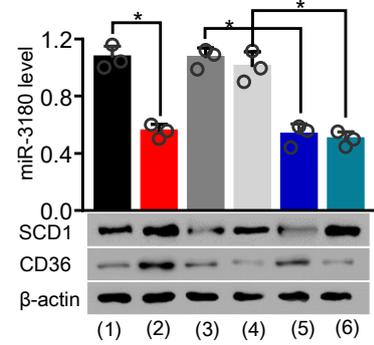
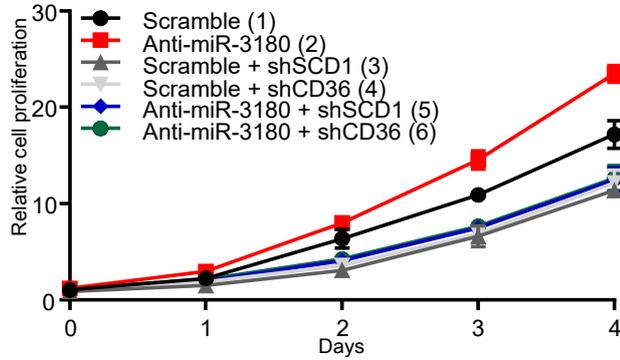
B



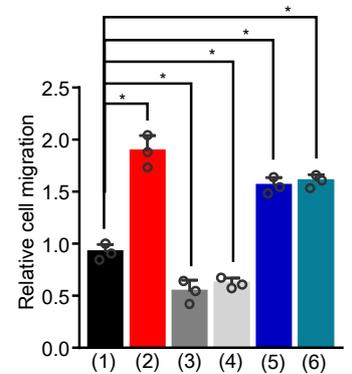
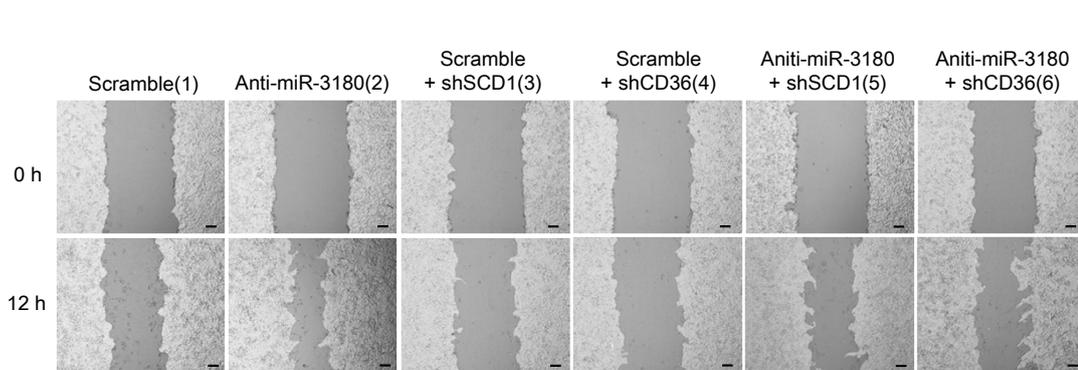
C



D



E



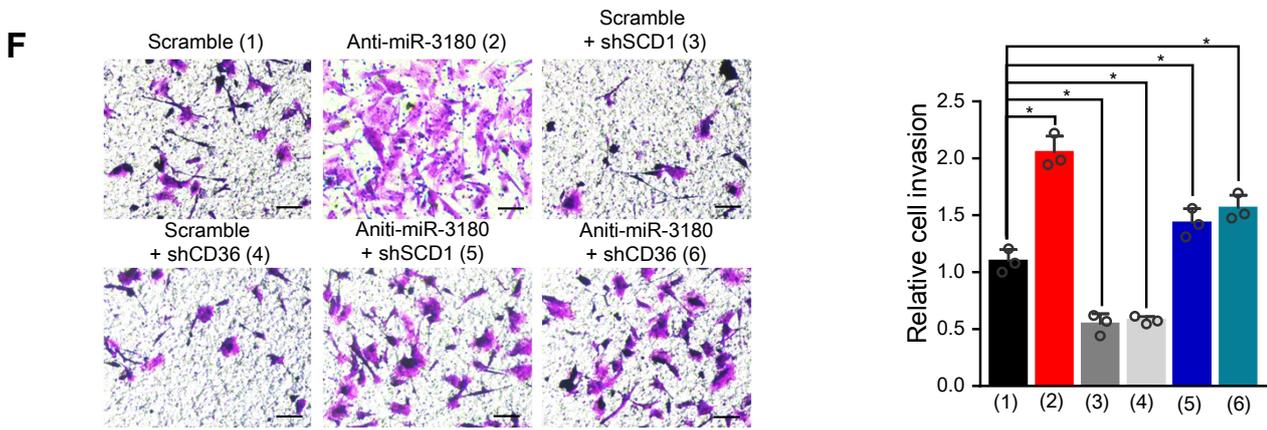
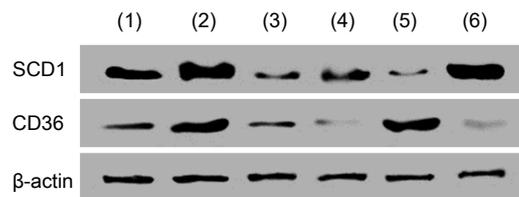
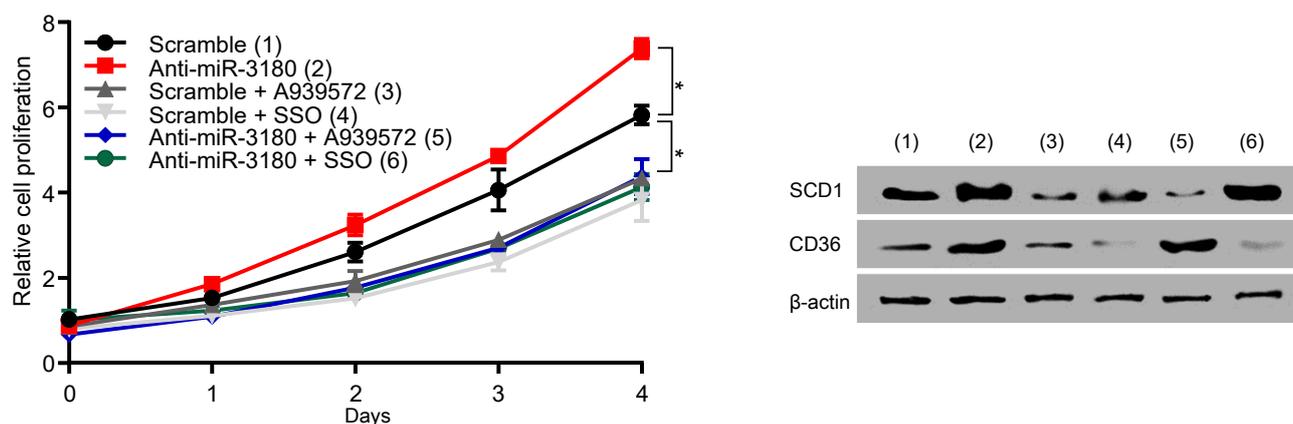


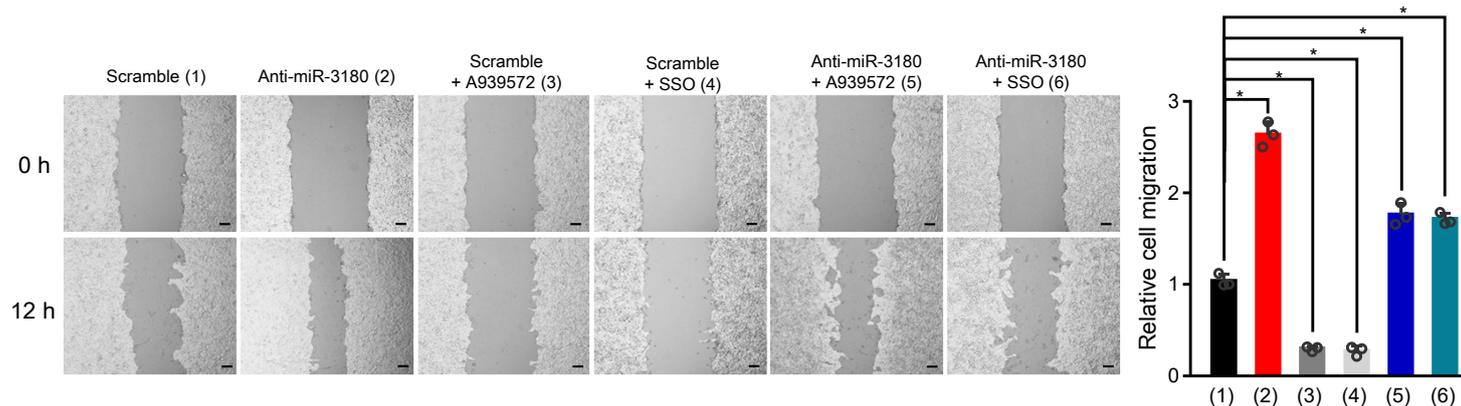
Figure S2. MiR-3180 inhibits the proliferation, migration, and invasion of HCC cells in CD36- and SCD1-dependent manner. (A) Cell proliferation analysis of MHCC-97H cells transfected with indicated expression vectors. The expression of miR-3180, CD36 and SCD1 are demonstrated in the right panel. (B) Wound healing analysis of MHCC-97H cells transfected as (A). Scale bar: 50 μ m. Statistical analysis of relative migrations are demonstrated shown in the right panel. (C) Transwell analysis of MHCC-97H cells transfected as (A). Scale bar: 20 μ m. Statistical analysis of invasion cells are shown in the right panel. (D) Cell proliferation analysis of MHCC-97H cells transfected with indicated expression vectors. The expression of miR-3180, CD36 and SCD1 are demonstrated in the right panel. (E) Wound healing analysis of MHCC-97H cells transfected as in (D). Scale bar: 50 μ m. Statistical analysis of relative migrations are demonstrated shown in the right panel. (F) Transwell analysis of MHCC-97H cells transfected as in (D). Scale bar: 20 μ m. Statistical analysis of invasion cells are shown in the right panel. Data shown are presented as mean \pm SD of triplicate measurements with similar results. **P < 0.01.

Figure S3

A



B



C

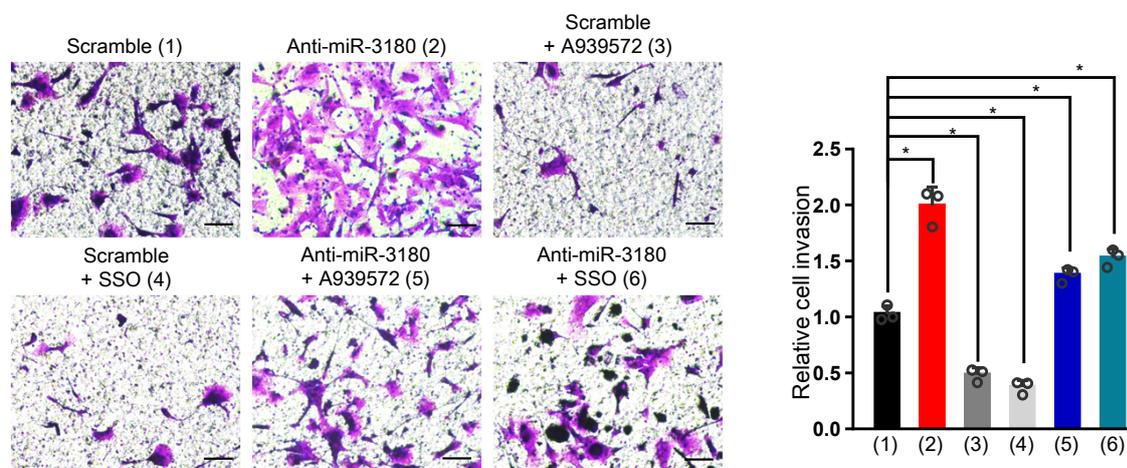


Figure S3. MiR-3180 inhibits the proliferation, migration, and invasion of HCC cells by suppressing lipid synthesis and uptake. (A) Cell proliferation analysis of MHCC-97H cells transfected with indicated expression vectors and treated with or without A939572 (15 μ m) or SSO (50 μ m). The expression of CD36 and SCD1 are shown in the right panel. (B) Wound healing analysis of MHCC-97H cells transfected and treated as (A). Scale bar: 50 μ m. Statistical analysis of relative migrations are shown in the right panel. (C) Transwell analysis of MHCC-97H cells transfected and treated as (A). Statistical analysis of relative invasion are shown in the right panel. Scale bar: 20 μ m. Data shown are presented as mean \pm SD of triplicate measurements with similar results. **P < 0.01.

Figure S4

A

- Scramble + control shRNA (1)
- Anti-miR-3180 + control shRNA (2)
- Scramble + SCD1 shRNA (3)
- Scramble + CD36 shRNA (4)
- Anti-miR-3180 + SCD1 shRNA (5)
- Anti-miR-3180 + CD36 shRNA (6)

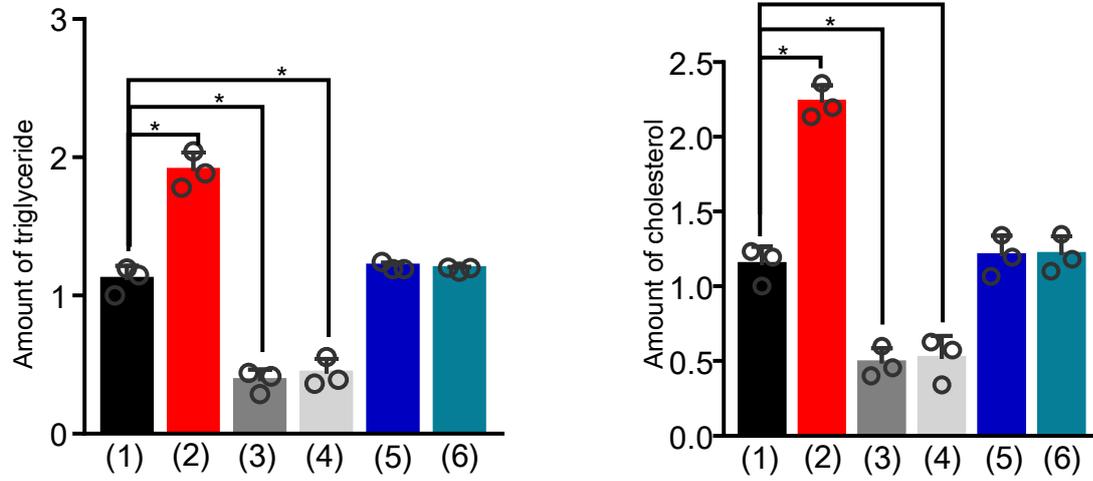


Figure S4 The miR-3180 inhibits lipid synthesis *in vivo*. Relative triglyceride and cholesterol contents in the indicated xenograft tumors from Figure 7A,B are shown.