Supplemental Figure 7



Supplemental Figure 7.

A and B, Liquid chromatography-tandem mass spectrometry (LC/MS) analysis assessing free cholesterol and cholesterol ester species from A498 cells subcutaneously implanted in nude mice treated or not with BLT-1 (50 mg/kg) by oral gavage daily for 30 days. **C**, AKT phosphorylation, AKT, SCARB1 and PDZK1 protein expression assessed by immunoblots in shSCR and shPDZK1 A498 cells grown in media supplemented with 10% FBS. GAPDH was used as the loading control. D, ROS levels assessed by flow cytometry measuring DCFDA fluorescence in A498 cells cultured in 10% FBS, 10% DLPS or 10% DLPS supplemented with N-acetyl-L-cysteine (NAC) (10mM) media for 72h. Representative plots (left) and mean fluorescence intensity quantifications are shown (right). E, Representative photographs of A498 cells grown in media supplemented with 10% FBS or 10% DLPS and treated with or without N-acetyl-L-cysteine (NAC) (10mM) for 72h. Magnification (100X) F. AKT phosphorylation and AKT protein expression assessed by immunoblots in A498 cells grown in media supplemented with 10% FBS or 10% DLPS and treated with or without N-acetyl-L-cysteine (NAC) (10mM) for 72h. GAPDH was used as the loading control. G, ROS levels assessed by flow cytometry measuring DCFDA fluorescence in shSCR and shSCARB1 A498 cells cultured in 10% FBS, 10% DLPS or 10% DLPS supplemented with N-acetyl-L-cysteine (NAC) (10mM) media for 72h. Representative plots (left) and mean fluorescence intensity quantifications are shown (right). H, ROS levels assessed by flow cytometry measuring DCFDA fluorescence in shSCR and shSCARB1 A498 cells cultured in 10% FBS, 10% DLPS or 10% DLPS supplemented with a-tocopherol (0.5mM) media for 72h. Representative plots (left) and mean fluorescence intensity quantifications are shown (right). (All experiments were performed in at least triplicates and statistical analysis was applied with *=P<0.05, **=P<0.01, ***=<0.001, n.s=non-significant).