Supplemental Figure 4



Supplemental Figure 4.

A, Representative photographs of immunohistochemistry analysis performed on BIOMAX tissue microarrays for SCARB1 expression in normal kidney tissue and ccRCC tumors. Magnification (100X). B, Quantification of immunohistochemistry analysis performed in A. Normal n=40, ccRCC n=100. C, Representation of SCARB1 mRNA expression across RCC subtypes analyzed using cBioPortal datasets. Chromophobe n=197, ccRCC n=1461, Papillary n=574. D, SCARB1 protein expression in normal kidney tissue, ccRCC tumor, normal kidney cell lines and ccRCC cell lines assessed by immunoblots. GAPDH was used as the loading control. E, Representative photographs of shSCR and shSCARB1 HK-2 cells grown in media supplemented with 10% FBS. Magnification x100. F, SCARB1 protein expression assessed by immunoblots in shSCR and shSCARB1 HK-2 cells. GAPDH was used as the loading control. G, Annexin-V/PI staining and flow cytometry analysis performed on shSCR and shSCARB1 HK-2 cells. H, Representative photographs of HK-2 cells grown in media with 10% FBS and treated with BLT-1 (5mM) or vehicle control (DMSO). Magnification x100. I, Annexin-V/PI staining and flow cytometry analysis performed on HK-2 cells grown in media with 10% FBS and treated with BLT-1 (5mM) or vehicle control (DMSO). J, Representative photographs of shSCR and shSCARB1 RPTEC cells grown in media supplemented with 10% FBS. Magnification x100. K, Annexin-V/PI staining and flow cytometry analysis performed on shSCR and shSCARB1 RPTEC cells. L, Representative photographs of RPTEC cells grown in media with 10% FBS and treated with BLT-1 (5mM) or vehicle control (DMSO). Magnification x100. M, Annexin-V/PI staining and flow cytometry analysis performed on RPTEC cells grown in media with 10% FBS and treated with BLT-1 (5mM) or vehicle control (DMSO).