

SUPPLEMENTARY FIGURES AND FIGURE LEGENDS

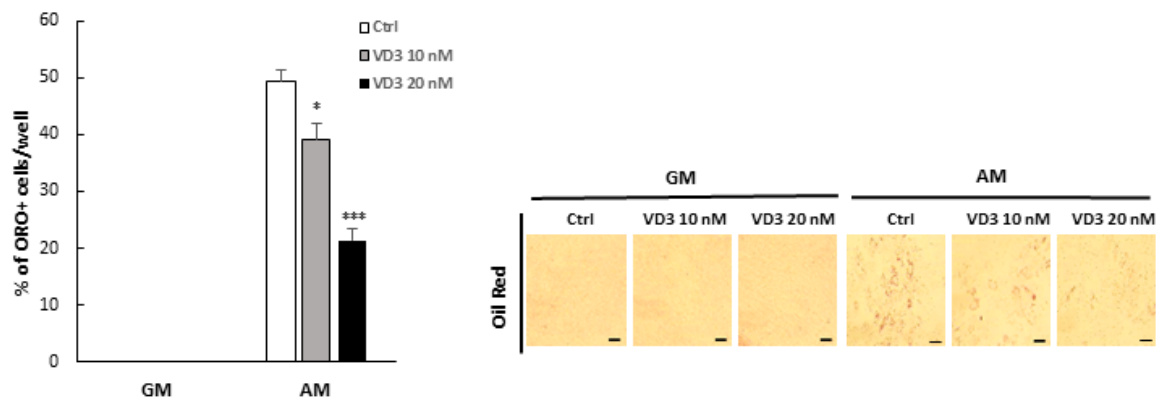


Figure S1. VD3 modulates adipogenic differentiation of BM-MSCs. Adipogenesis of BM-MSCs was analyzed for cells grown in GM or adipogenic medium (AM) with or without VD3 (10nM or 20nM). After 28 days intracellular lipid droplets were detected by Oil Red O staining. Representative images are shown (Scale bars: 20µm). Graph illustrates quantitative analysis showing Oil Red O positive (ORO+) cells percentage enumerated per visual field. Results are presented as means \pm SEM for at least three independent experiments. Statistically significant differences by t test: * $p < 0.05$; *** $p < 0.001$ compared to untreated AM control.

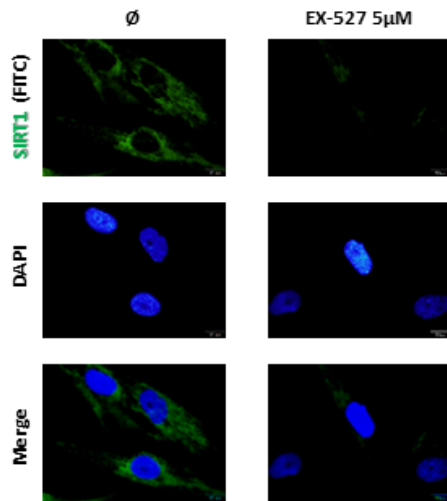


Figure S2. EX-527 inhibits SIRT1 expression in BM-MSCs. BM-MSCs were treated with or without EX-527 (5µM) during 5 days under standard cultivation conditions in GM. Protein expression of SIRT1 following the treatment detected by indirect immunofluorescence staining. Secondary fluorescein isothiocyanate (FITC)-conjugated antibodies were used for labeling corresponding primary antibodies. DNA was stained with 4',6-diamidino-2-phenylindole (DAPI). Representative images from two independent experiments are shown (Scale bars: 20µM).

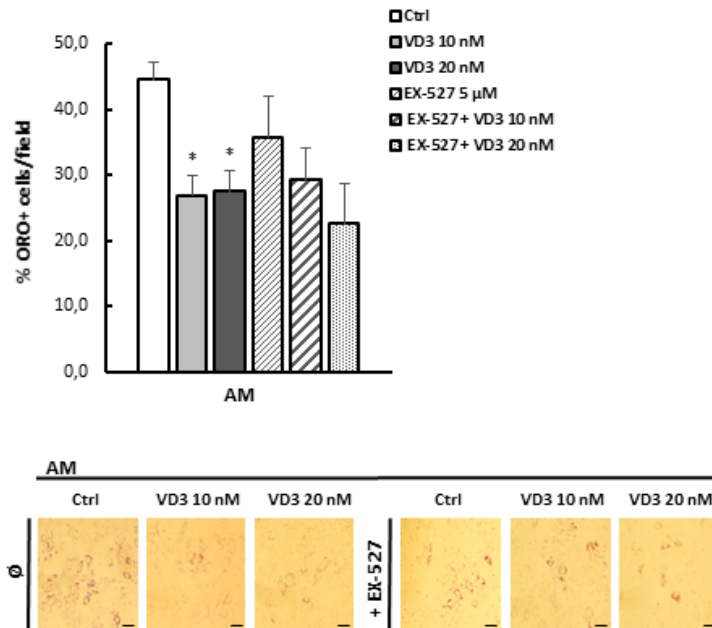


Figure S3. Reduced adipogenic differentiation of VD3-pretreated BM-MSCs: role of SIRT1 signaling. BM-MSCs were pretreated for 5 days with VD3 (10nM or 20nM), EX-527 (5 μ M) or both in GM. Afterwards, cells were cultured in AM for appropriate period to induce adipogenesis. Intracellular lipid droplets were detected after 28 days cultivation in AM by Oil Red O staining. Representative images are shown (Scale bars: 20 μ m). Graphs represent quantitative analysis of Oil Red O positive (ORO+) cells percentage enumerated per visual field. Results are presented as means \pm SEM for at least three independent experiments. Statistically significant differences by t test: * $p < 0.05$ compared to untreated AM control.