Figure S1. Effect of madanin-1 WT and madanin-1 2S on the viability of (A) SKOV3 and (B) MDA-MB-231 cells. The SKOV3 and MDA-MB-231 cells were treated for 48 h with or without thrombin (2 U/ml) after pretreatment with madanin-1 WT or madanin-1 2S at different concentrations (0–20 µg/ml) for 1 h. Cell viability was assessed with an MTT assay. The results showed that no significant changes in cell viability were induced in SKOV3 and MDA-MB-231 cells by treatment with madanin-1 WT or madanin-1 2S. Assays were performed in triplicate using cell lines (n=3). Data are expressed as percentages of control values and are presented as the mean ± standard deviation. WT, wild-type; 2S, 2 sulfation.
Figure S2. Effect of madanin-1 WT and madanin-1 2S on the apoptosis of SKOV3 cells. The SKOV3 cells were treated for 48 h with or without thrombin (2 U/ml) after pretreatment with (A) madanin-1 WT or (B) madanin-1 2S at different concentrations (0-20 µg/ml) for 1 h. Apoptosis of SKOV3 cells was analyzed by flow cytometry. The results showed that no significant level of apoptosis was induced in SKOV3 cells by treatment with madanin-1 WT or madanin-1 2S. The percentage of cells is indicated in each quadrant. WT, wild-type; 2S, 2 sulfation.
Figure S3. Effect of madanin-1 WT and madanin-1 2S on the apoptosis of MDA-MB-231 cells. The MDA-MB-231 cells were treated for 48 h with or without thrombin (2 U/ml) after pretreatment with (A) madanin-1 WT or (B) madanin-1 2S at different concentrations (0-20 µg/ml) for 1 h. Apoptosis of MDA-MB-231 cells was analyzed by flow cytometry. The results showed that no significant level of apoptosis was induced in MDA-MB-231 cells by treatment with madanin-1 WT or madanin-1 2S. The percentage of cells is indicated in each quadrant. WT, wild-type; 2S, 2 sulfation.