



Supplementary Figure 1. Conditioned media from rDrm-treated SIE cells were added for 10 minutes to naïve serum-starved SIE cells in the absence or in the presence of neutralizing anti-Ang-1 antibodies. Then, immunoprecipitation with anti-Tie-2 antibodies was performed on the cell extracts followed by Western blotting with anti-phosphotyrosine antibodies (Santa Cruz). Uniform loading of the gel was confirmed by incubation of the membrane with anti-Tie-2 antibodies.