

Preclinical efficacy of maternal embryonic leucine-zipper kinase (MELK) inhibition in acute myeloid leukemia

Supplementary Material

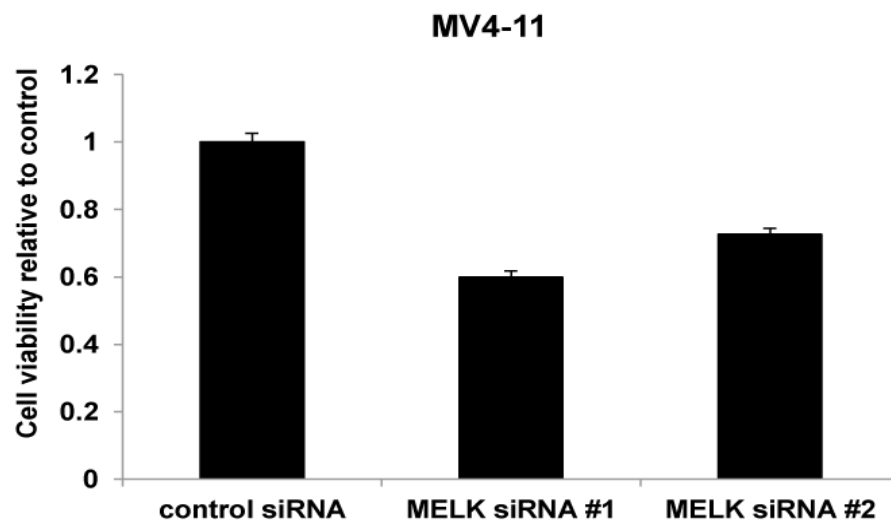


Figure S1: MELK knock-down decreased cell viability. MV4-11 cell transfected with two different siRNAs against MELK and cell viability was assessed by MTT assay 48 hours later.

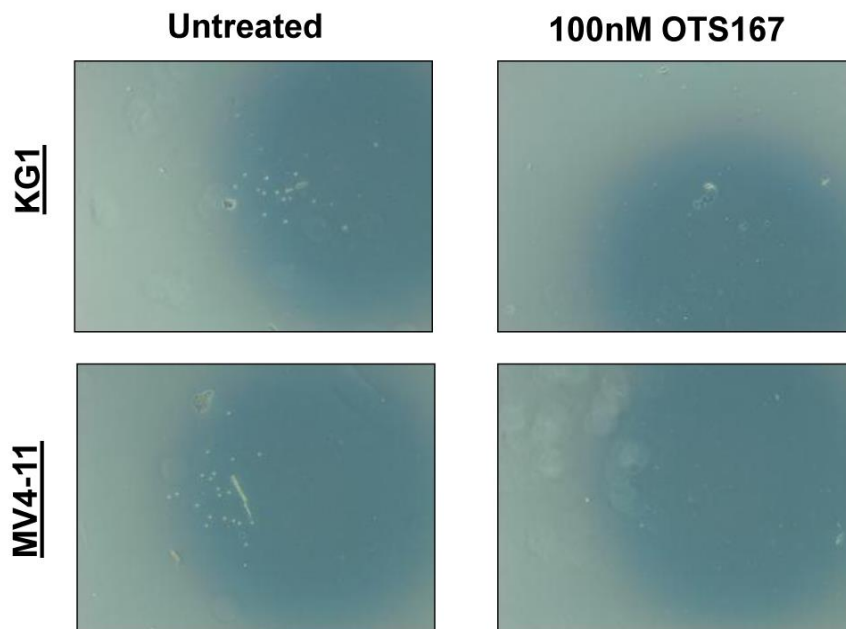


Figure S2: MELK inhibitor OTS167 reduces cell migration. KG1 and MV4-11 cells were treated with 100nM of OTS167 for 6 hours, then cell migration was assessed in trans-well plate, image shows cells in the bottom well.

(12 hours)

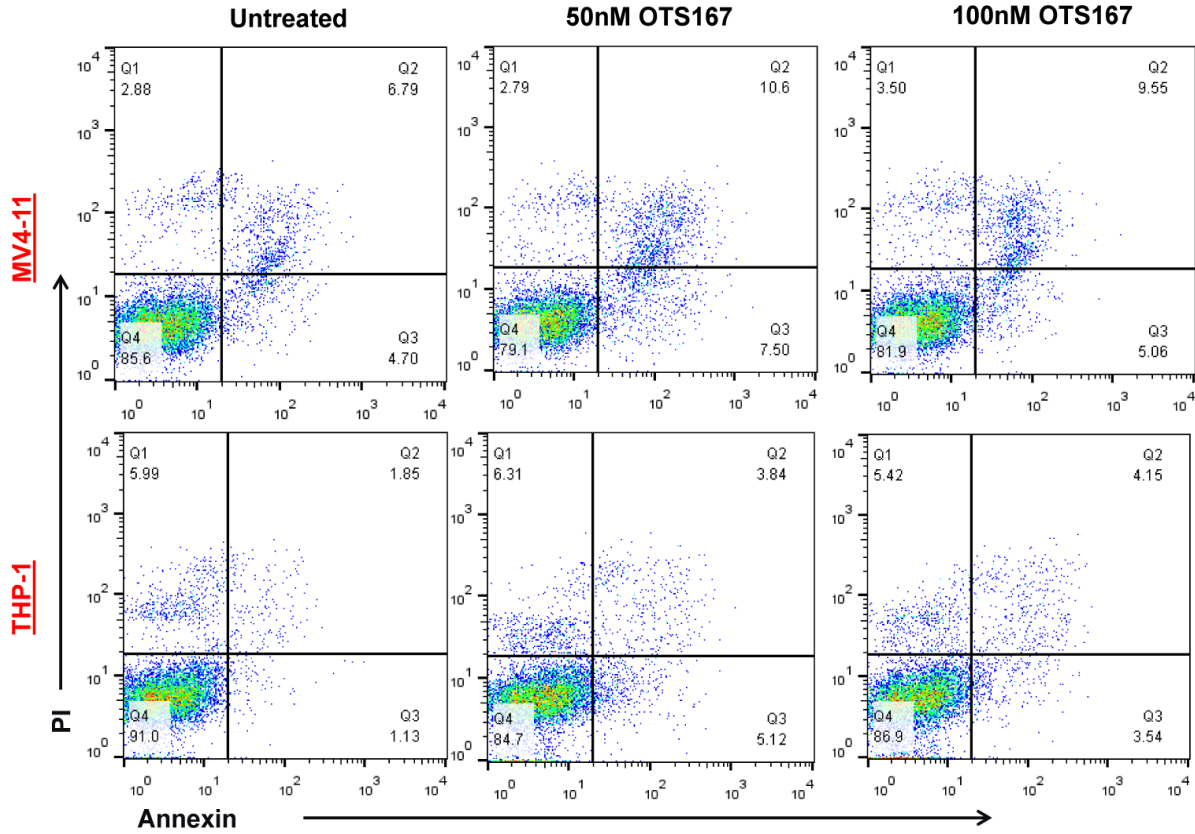


Figure S3: The effect of MELK inhibitor OTS167 on cell apoptosis 12 hours following treatment. MV4-11 and THP-1 cells were treated with 50 and 100nM of OTS167 for 12 hours, then apoptosis was assessed by Annexin and PI staining.

24 hours

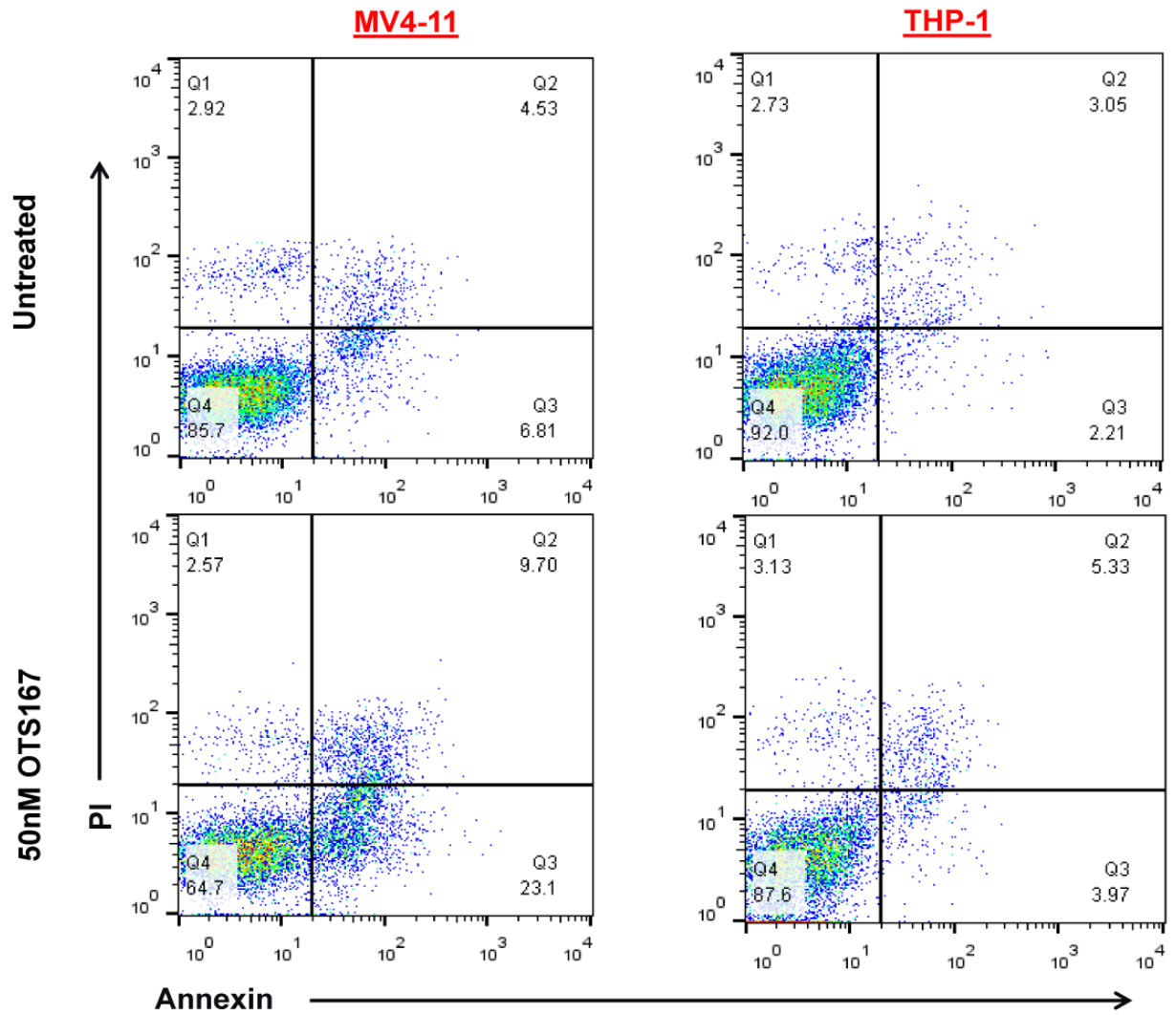


Figure S4: The effect of MELK inhibitor OTS167 on cell apoptosis 24 hours following treatment. MV4-11 and THP-1 cells were treated with 50nM OTS167 for 24 hours, then apoptosis was assessed by Annexin and PI staining.

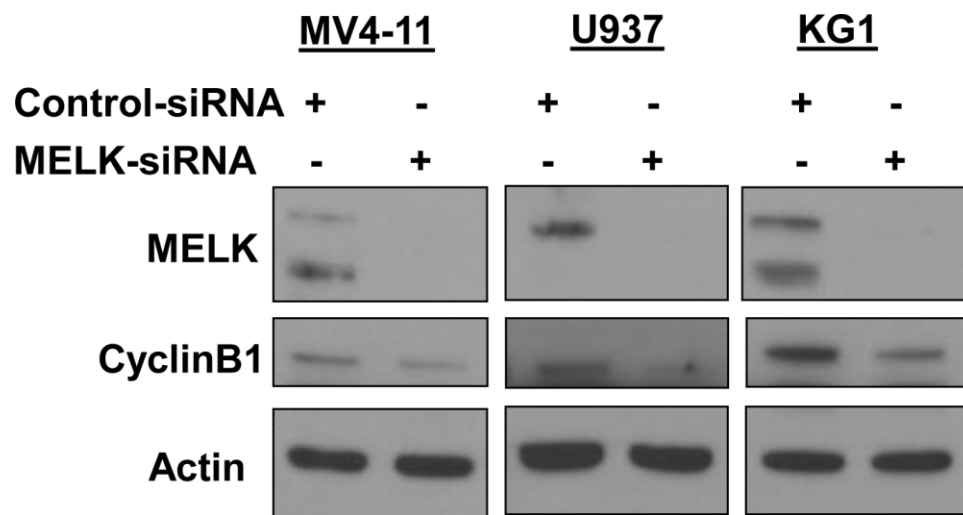


Figure S5: The effect of MELK inhibition on CyclinB1 expression. MV4-11, U937 and KG1 cells were transfected with MELK-siRNA, and cyclinB1 expression was assessed 48 hours following transfection.