Figure S6. SpiD3 inhibits NF-κB activity and diminishes CD40L-induced survival signaling in CLL.

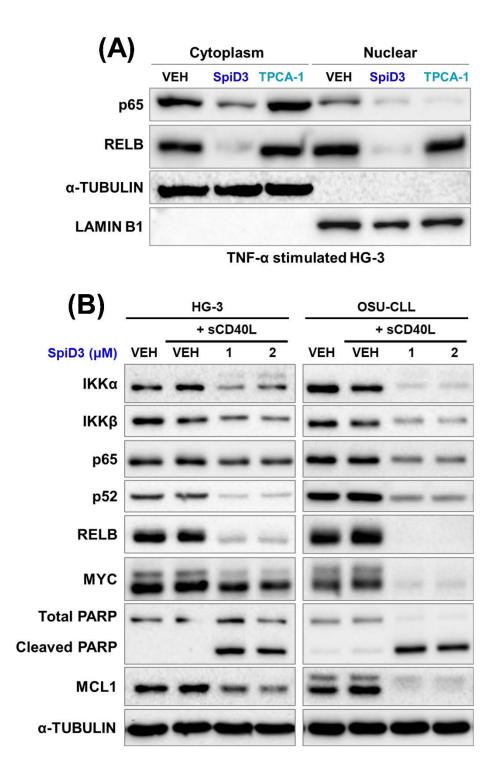


Figure S6. SpiD3 inhibits NF-κB activity and diminishes CD40L-induced survival signaling in CLL.

(A) HG-3 cells were treated with SpiD3 (2 μ M), TPCA-1 (10 μ M), or equivalent DMSO vehicle (VEH) for 4 h and stimulated with TNF- α (20 ng/mL) during the last 15 min of treatment (n = 4 independent experiments). Cytoplasmic and nuclear fractions were subjected to immunoblotting probing for p65 and RELB. α -TUBULIN served as the cytoplasmic fraction loading control and LAMIN B1 served as the nuclear fraction loading control. (B) Immunoblot analysis of the indicated proteins in whole cell lysates of HG-3 and OSU-CLL cells treated with SpiD3 (1, 2 μ M) for 4 h and co-currently stimulated with sCD40L (500 ng/mL). α -TUBULIN served as the loading control (n = 4 independent experiments/cell line).