Figure S11. FK866-mediated NAD⁺ depletion mediates FK866’s cytotoxic activity. A, Primary B-CLL and AML cells were incubated in 24-well plates in the presence or absence of 10 nM FK866, 100 μg/ml VA, 500 μM BU, or their combination. 48 h later, NAD⁺ levels were determined by enzymatic cycling assay. NAD⁺ values were normalized to protein concentrations (expressed in mg). B, Primary AML cells were plated in 96-well plates and incubated with 100 nM FK866. NAD⁺ was added to the medium every 12 h in order to achieve the indicated concentrations. Viability was quantified after 96 h of incubation by PI staining and flow-cytometry.