Supplementary Figures

Transcriptional mechanism of vascular endothelial growth factor-induced expression of protein kinase CβII in chronic lymphocytic leukaemia cells.

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Supplementary Figure 1 <u>Treatment of Mec1 cells with mithramycin or SP1-specific siRNA reduces PKC6 mRNA levels</u>. 2x10⁶ MEC1 were either were cultured for 24h in the presence of the indicated concentrations of mithramycin (in nmol/L), or were nucleofected with the indicated SP1-specific (Oligo 1, 2, 3 and mix) or control siRNA (NEG) oligonucleotides and then cultured as described in the materials and methods for each cell type. Harvested cells were were analyzed by qRT-PCR for SP1 (**A**, **B**) or for PKCβII mRNA (**C**) using RNApolII as a reference gene. The results presented represent the mean±SE of n=3 separate replicates. Statistical analysis for all parts in this figure were performed using a students t-test for paired data. UT - untreated



Supplementary Figure 2 <u>Analysis of STAT3 repression of PRKCB promoter function in Mec1 cells</u>. 2x10⁶ MEC1 were cotransfected with (0.1µg) pRL and (2µg each) of wt pGL3-pkcβ-1.2kb, or site 1, site 2+3, or site 4 mutants of STAT3 binding sites within the *PRKCB* promoter of the pGL3-pkcβ-1.2kb construct. Luciferase assays were performed following 72h culture of the cells under serum-rich conditions, and are reported relative to renilla expression. The data represented mean±SE of three independent experiments. Statistical analysis was performed using a student's t-test for paired data.



Supplementary Figure 3. Uncropped Western blot for SP1 represented in manuscript Figure 1E upper panel.



Supplementary Figure 4. Uncropped Western blot for PKC β II represented in manuscript Figure 1E middle panel.



β-actin

Supplementary Figure 5. Uncropped Western blot for β -actin represented in manuscript Figure 1E lower panel.

