

**Suppl. Tab. 1. Characteristics of AML cell lines.**

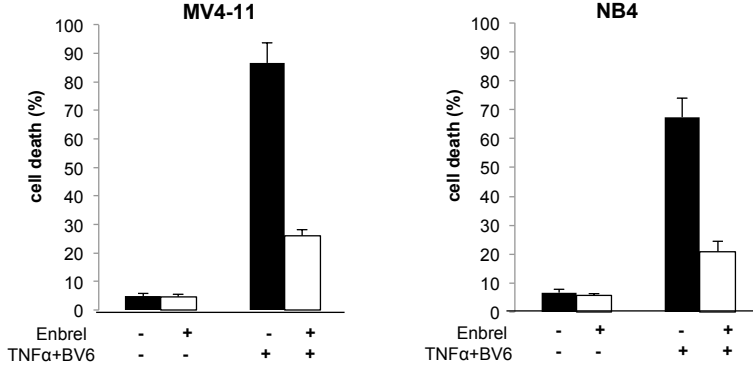
<b>Cell line</b>	<b>FAB classification</b>	<b>Genetics</b>
NB4	FAB M3	t(15;17) (q22;q11-12)
MV4-11	FAB M5	t(4;11)(q21;q23) FLT3-ITD mutation MLL-AF4 fusion
Molm13	FAB M5a	FLT3-ITD mutation MLL-AF9 fusion ins(11;9)(q23;p22p23)
MonoMac6	FAB M5	t(9;11)(p22;q23) MLL-AF9 fusion
OCI-AML3	FAB M4	NPM1-gene-mutation (Type A) DNMT3a R882C mutation

**Suppl. Tab. 2. Synergistic induction of apoptosis by BV6 and demethylating agents in AML.**

Cell line	BV6 + 5AC		BV6 + DAC	
	<i>CI</i>	<i>fa (%)</i>	<i>CI</i>	<i>fa (%)</i>
MV4-11	0.094	37.3	0.150	49.9
NB4	0.233	50.9	0.254	67.7
Molm13	0.271	56.6	0.161	64.3
MonoMac6	0.105	56.8	0.019	49.7
OCI-AML3	0.436	68	1.113	11.3

AML cell lines were treated for 72 hours with BV6 and/or 5AC or DAC as indicated in the figure legend of Fig. 1. Apoptosis was determined by forward/side scatter analysis and flow cytometry. Combination index (CI) and fraction affected (fa) were calculated by CalcuSyn software as described in Materials and Methods. CI values and fraction affected are indicated for the following concentrations: MV4-11: 300 nM 5AC/100 nM BV6, 50 nM DAC/100 nM BV6; NB4: 300 nM 5AC/600 nM BV6, 30 nM DAC/600 nM BV6; Molm13: 300 nM 5AC/100 nM BV6, 50 nM DAC/100 nM BV6; MonoMac6: 600 nM 5AC/3  $\mu$ M BV6, 300 nM DAC/3  $\mu$ M BV6; OCI-AML3: 100  $\mu$ M 5-AC/2  $\mu$ M BV6, 6  $\mu$ M DAC, 2  $\mu$ M BV6. CI <0.9 indicates synergism, 0.9-1.1 additivity and >1.1 antagonism.

Suppl. Fig. 1



## **Supplementary Figure legend**

### **Supplementary Fig. 1. Enbrel blocks BV6/TNF $\alpha$ -induced cell death.**

MV4-11 and NB4 cells were treated for 48 hours with BV6 (MV4-11: 600 nM BV6, NB4: 100 nM BV6) and/or 30 ng/ml TNF $\alpha$  in the presence or absence of 100  $\mu$ g/ml Enbrel. Cell death was determined by forward/side scatter analysis. Mean and SD of three experiments performed in triplicate are shown.