



Supplemental Material to:

**Kenichiro Doi, Qiang Liu, Krishne Gowda, Brian M Barth,
David Claxton, Shantu Amin, Thomas P Loughran Jr,
and Hong-Gang Wang**

**Maritoclax induces apoptosis in acute myeloid leukemia
cells with elevated Mcl-1 expression**

**Cancer Biology & Therapy 2014; 15(8)
<http://dx.doi.org/10.4161/cbt.29186>**

<http://www.landesbioscience.com/journals/cbt/article/29186/>

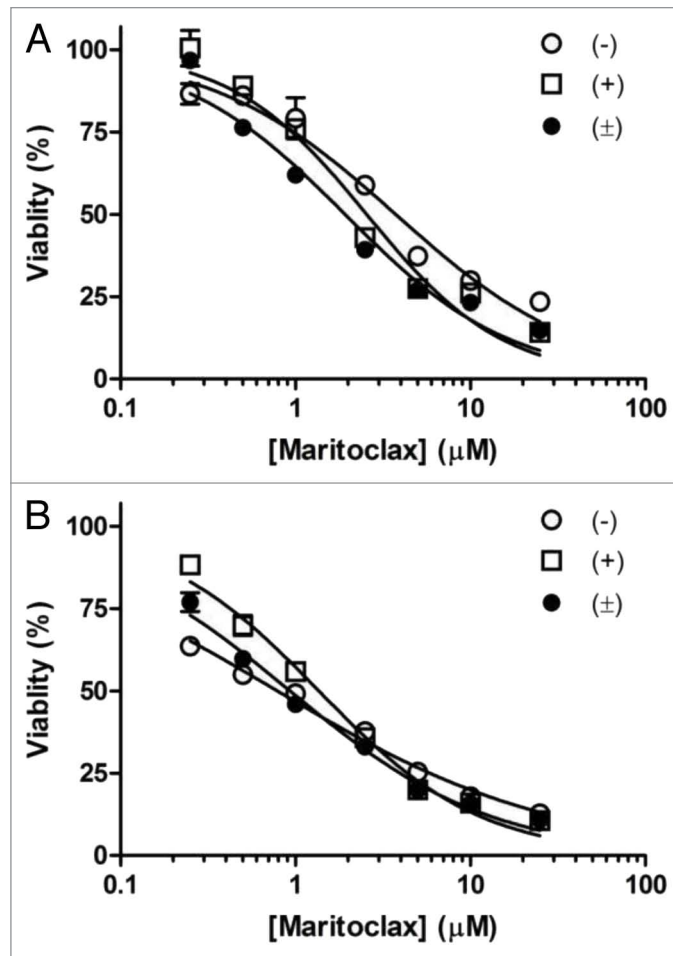


Figure S1. Potency of (+), (-), and racemic maritoclax in U937 (A) and Mcl-1-IRES-Bim K562 cells (B) in viability assays.

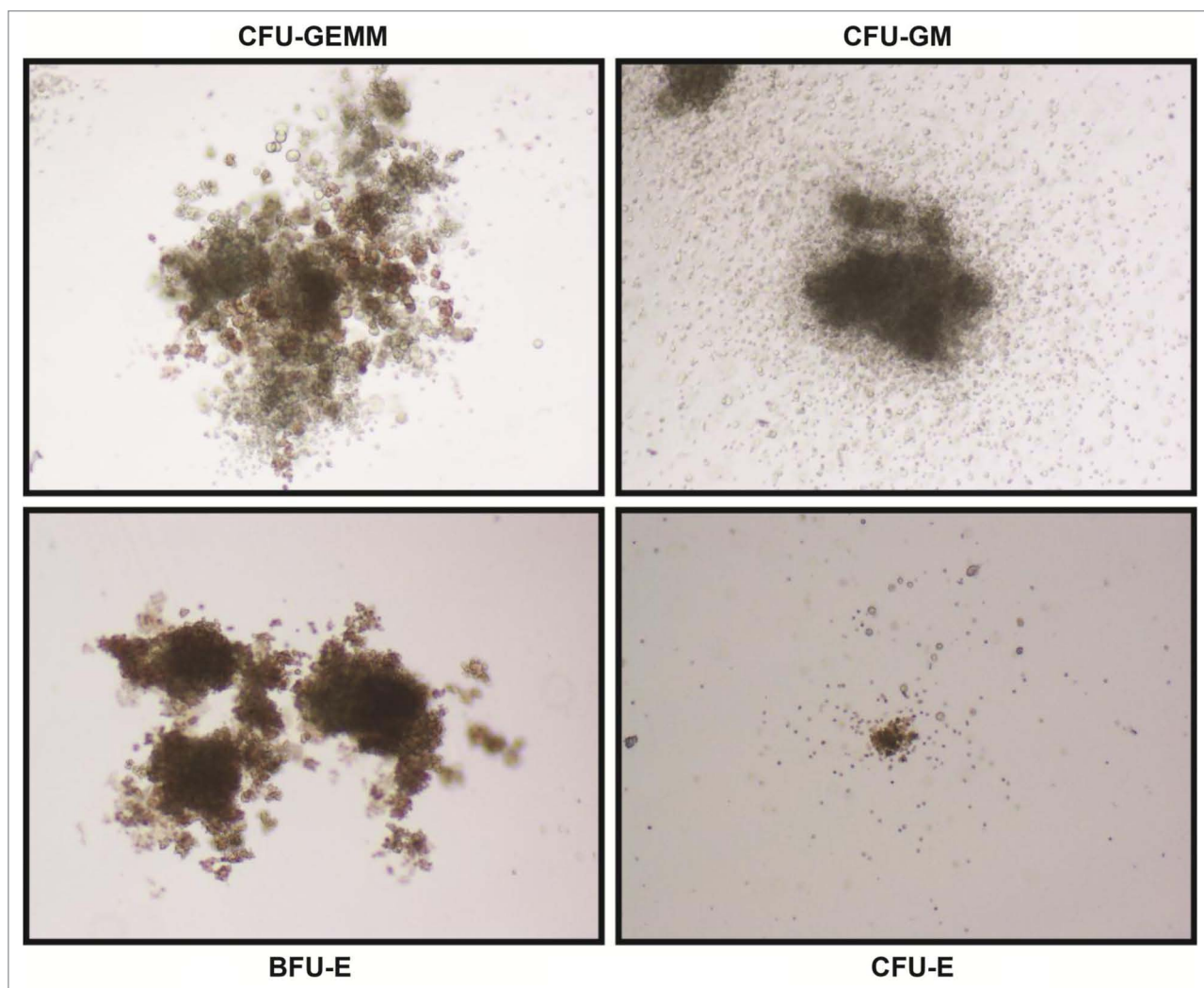


Figure S2. Representative colonies in vehicle-treated colony formation assay of primary mouse bone marrow, taken with the Olympus DP20 and the cellSens Standard software at 10× magnification.

Table S1. Primary human AML sample characteristics

Patient Case #	WBC ($\times 1000/\mu\text{L}$)	Blast (%)	NPM1 Mutation	FLT3-ITD Mutation	Cytogenetics
555	15.78	70	ND	ND	t(9;11)
559	66.6	66	-	-	Complex
477	122.3	78	-	-	del(15)(q13q15)
574	107.7	58	+	+	Normal

Table S2. The EC₅₀ of MEF, HEK293, and HeLa cells to maritoclax in viability assays

	EC ₅₀ (μM)		
	Maritoclax	ABT-737	Daunorubicin
MEF	18.1*	> 50*	1.75*
HEK293	> 50*	> 50*	41.2*
HeLa	> 50	49.4	0.33

*Cells were treated with the indicated compounds for 24 h.