Class	Name	EBV	Reference
HL	L428 (cHL)	-	[1,2]
	KM-H2 (cHL)	-	[2]
	HDLM2 (T-cell HL)	-	[2]
	L1236 (cHL)	-	[2,3]
	L591 (cHL)	+	[2]
	DEV (NLPHL)	-	[4,5]
PMBL	MedB-1	-	[6]
	Karpas 1106P	-	[6]
DLBCL	ROSE	-	[2]
	VER	-	[2]
	SU-DHL-4	-	A. Epstein (UCLA)
	SU-DHL-6	-	A. Epstein (UCLA)
	OCI Ly3	-	[7]
	SCHI	-	
BL	Raji	+	[8]
	CA46	-	[8]
	BL-65	+	[8]
	Namalwa	+	[8]
	DG-75	-	[8]
	Jiyoye	+	[8]
	Ramos (RA 1)	-	[8]
LCL	CB-LCL 6.28	+	[9]
	CB-LCL 5.5-1	+	[9]
	CB-LCL 5.b2	+	[9]
	CB-LCL 6.16	+	[9]
	CB-LCL 5.B8	+	[9]
	1H2 (EBV transformed)	+	[10]
	15E6 (EBV transformed)	+	[10]
CLL	EHEB (B-CLL)	+	DSMZ
	MEC-1 (B-CLL)	+	DSMZ
	MEC-2 (B-CLL)	+	DSMZ
	JVM-3 (B-CLL)	+	DSMZ

Table W1. Cell Lines Used for miRNA Expression Validation.

DSMZ = Deutsche Sammlung von Microorganismen un Zellkulturen, GmbH, located in Braunschweig, Germany.

Table W2. Primer Sequences Used for Amplification of Target Sequences.

Gene	Forward Primer* (5'-3')	Reverse Primer [†] (5'-3')	Size (bp)
GAB2	GAGCTC-ATGTCAGATCGCAGGGTAGG	TCTAGA-AGTCGCTCTCCGAAGATTCC	1205
BACH2	GAGCTC-GGAAAGACAGCAGTGATGAC	GCGGCCGC-GTGCAAGTGGCAAAGTTGAC	659
ICOSL	GAGCTC-AGGGCCGTGTTTGGCTACAG	TCTAGA-CCTCAGGCATGAGGGACAGA	1187
ZIC3	GAGCTC-TGGTACGTCTGAGGACAAAC	GCGGCCGC-GAGTCTTCCCAGATGGAAAC	979
IKBKE	GAGCTC-TCCCAGCACCTCCTGATGTC	GCGGCCGC-CTGCTTCCCAGGGAGAAAGG	215
TIP120	TGTTTGGCTTTCTTCCATTG	AGAAAGAAATTCATGGTCAC	226
AGTR1	GAGCTC-CATGTTCGAAACCTGTCCATAAAG	GCGGCCGC-ATAAAATTATTTTATTTTAAAGTAAAT	897
ZNF537	GAGCTC-TGTGGAAGGCACCTTCAG	TCTAGA-CCAGCTCCGTCATAAACAG	1668
FGF7	ACTACTCGAGCTGATCAAGCTGGACTTGCGC	ATAAGAATGCGGCCGCTAACAAAACAATAAAATTCAAATACAAC	790
MNAB	TTGTGACCACACCATGGAAG	TTACTGCACCGGATTGCTAC	282
MAF	GAGCTC-AAGCCTGCATCAACCTTCTG	GCGGCCGC-GGGCAATTATGGCTCAACTC	1795

TIP120, MNAB, and FGF7 were subcloned first in a TOPO vector and lack specific restriction sites. Size indicates the size of the PCR product that was cloned into the vector. *The restriction site for endonucleases *Not*I was added to the forward primer. [†]The restriction site for *Sst*I or *Xba*I was added to the reverse primer (separated by dash).



Figure W1. Heat chart of miRNA expression in B-cell–derived cell lines.



Figure W2. Schematic presentation of the predicted target sequences cloned in the psiCHECK2 vector. The regions from the 3' UTR of potential miR-155 targets (A) were cloned into psiCHECK2 vector (B) for target gene validation experiments.



Figure W3. The FL/RL luciferase expression ratios observed in the three HL cell lines without miR-155–specific inhibitor. Luciferase ratios obtained for the psiCHECK2 constructs containing 3' UTR sequences of potential miR-155 target genes. Error bars indicate the SD for three replicate experiments.