Contino et al., supplementary table S1

Table S1. List of gene-specific oligonucleotides used in this study

(**A**) Oligonucleotides used for ERBB2 promoter luciferase reporter construct		
Target sites*	Sense (S) and Antisense (A) sequences (5' to 3')[§]	Amplicon size (bp)
HER-5	S: GCTAGCGCTGGTCATGGTGGCACA	787
HER-6	S: GCTAGCACTTCAAAGATTCCAGAAGATATGC	558
HER-7	S: GCTAGCCACCAGCCTCTGCATTTAGG	306
HER-3	A: GAAGATCTGGGCTCCCCTGGTTTCTC	-
(**B**) Real time PCR oligonucleotides for ChIP assay		
Target genes	Forward (F) and reverse (R) sequences (5' to 3')	Amplicon size (bp)
ERBB2	ERP1-F: ACTTCAAAGATTCCAGAAGATATGC ERP2-R: GCTTGCATCCTACTCCATCC	162
ERBB2	ERP3-F: ACACATCCCCCTCCTTGACT ERP4-R: CGGAGAATCCCTAAATGCAG	228
ERBB2	ERP5-F: CTCTGCATTTAGGATTCTCCG ERP7-R: GGGCTCCCCTGGTTTCTC	294
MYC	MP3-F: AGGGCTTCTCAGAGGCTTG MP4-R: TGCCTCTCGCTGGAATTACT	113
MYC	MD-F: ATT GTC CCC TCT CCT CCT GT MD-R: CTT CGT CTC CCC TAC TGC TG	166

Oligonucleotide positions in the ERBB2 promoter sequence are shown in supplementary Figure 1S. [§] The underlined sequence refers to the NheI or BglII restriction site added to each oligonucleotide to clone the amplified fragment in the pGL3 reporter plasmid.