

Supplemental Table S1. PCR TOPO cloning Primer Pairs Sequences and Amplicon Sizes.

Supplemental Table 1

PCR TOPO Cloning Primer Pair Sequences and Amplicon Sizes

Gene	Sequence	Exon	Amplicon Size
KRAS	Forward-GTGAGTTGTATTAAAAGGTACTGG	2	265 bp
	Reverse-GGTCCCTGCACCAGTAATATGC		
KRAS	Forward-CCAGACTGTGTTCTCCCTTC	3	286 bp
	Reverse-TGCATGGCATTAGCAAAGAC		
PIK3CA	Forward-CTGTGAATCCAGAGGGAAA	9	269 bp
	Reverse-ACATGCTGAGATCAGCCAAA		
PIK3CA	Forward-CATTTGCTCCAAACTGACCA	20	389 bp
	Reverse-GGTCTTGCCTGCTGAGAGT		
TP53	Forward-CACTTGTGCCCTGACTTCA	5	267 bp
	Reverse-AACCAGCCCTGTCGTCTCT		
TP53	Forward-CTGCTCAGATAGCGATGGTG	6	251 bp
	Reverse-CTTAACCCCTCCTCCCAGAG		
TP53	Forward-CTTGGGCCTGTGTTATCTCC	7	199 bp
	Reverse-GGGTCAGAGGCAAGCAGA		
TP53	Forward-GGGAGTAGATGGAGCCTGGT	8	274 bp
	Reverse-TAACTGCACCCTTGGTCTCC		

Supplemental Table S2. Quantitative Real-Time PCR Primer Pairs. Amplicon sizes are within the range from 80 to 250 bp.

**Supplemental Table 2**  
**Quantitative Real-Time PCR Primer Pairs**

Gene	Sequence
ALDH1A1	CTCAAGGCCCTCAGATTGAC
	GTTTGGCCCTTCTTCTTC
OCT4	AGTGAGAGGCAACCTGGAGA
	ACACTCGGACCACATCCTTC
CD44	GGCGCAGATCGATTGAATA
	GAAAGCCTTGCAGAGGTAG
CD133	AATTCAACCAGCAACGAGTCC
	TCCAACAATCCATTCCCTGT
CD117	AAGTGGATGGCACCTGAAAG
	AGGGGCTGCTTCCTAAAGAG
CXCR4	TTGTGCCCTTAGCCCCTAC
	CACTTCCAATTCAAGCAAGCA
NANOG	CAGAAGGCCCTCAGCACCTAC
	ACTGGATGTTCTGGGTCTGG
RPL18	GGATGATCCGGAAGATGAAG
	CCGCACATCATCAGTTATGG

Supplemental Table S3. Summary of Identified Mutations.

**Supplemental Table 3**  
**Summary of Identified Mutations**

p53		Exon 5						Exon 6		Exon 7				Exon 8	
Control	CDS							703 c.703A>C AAC>CAC N235H Substitution							
	Codon														
100 mM	CDS	384 c.384T>C CCT>CCC P128P <b>Transition</b>	387 c.387C>T GCC>GCT A129A <b>Transition</b>	386 c.386C>T GCC>GTC A129V <b>Transition</b>	470 c.470T>G GTC>GGC V157G <b>Transversion</b>			590 c.590T>C GTG>GCG V197A <b>Transition</b>							
300 mM	CDS	384 c.384T>C CCT>CCC P128P <b>Transition</b>	388 c.388C>A CTC>ATC L130I <b>Transversion</b>	427 c.427G>C GTG>CTG V143L <b>Transversion</b>	436 c.436T>C TGG>CGG W146R <b>Transition</b>	488 c.488A>G TAC>TGC Y163C <b>Transition</b>	537 c.537T>C CAT>CAC H179H <b>Transition</b>	658 c.658T>G TAT>GAT Y220D <b>Transversion</b>	703 c.703A>C AAC>CAC N235H Substitution						
500 mM	CDS	384 c.384T>C CCT>CCC P128P <b>Transition</b>	386 c.386C>T GCC>GTC A129V <b>Transition</b>					658 c.658T>G TAT>GAT Y220D <b>Transversion</b>	688 c.688A>T ACC>AGC T230S Substitution	703 c.703A>C AAC>CAC N235H Substitution	710 c.710T>A ATG>AAG M237K Substitution	746 c.746G>C AGG>ACG R249T Substitution	751 c.751A>C ATC>ATA I251I Substitution	911 c.911C>A ACC>AAC T304N Substitution	914 c.914A>G AAA>AGA K305R Substitution

Transition	10
Transversion	5
Substitution	9

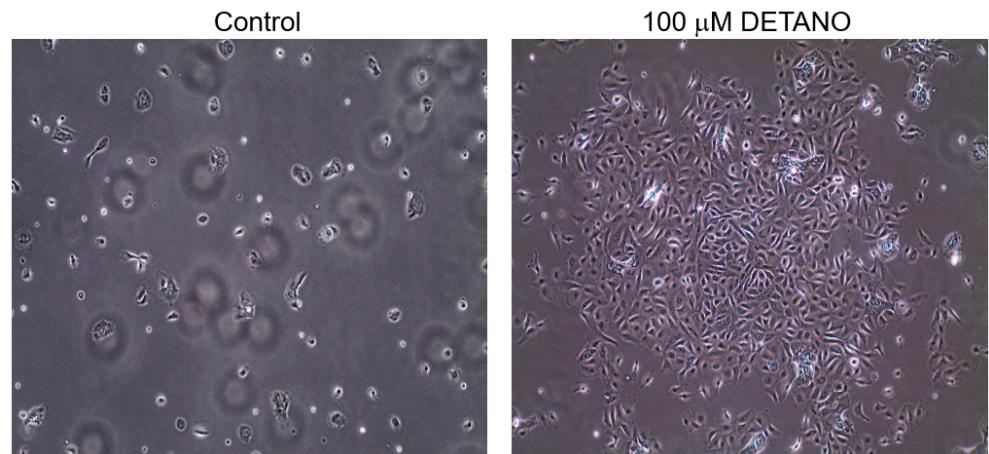
PIK3CA		Exon 9			Exon 20		
Control	CDS	1634 c.1634A>C GAG>GGC E545A <b>Transversion</b>	1658 c.1658delG AGT>ACC S553T Substitution	1659 c.1659T>C <b>Transition</b>	3088 c.3088A>G ACT>GCT T1030A <b>Transition</b>		
	Codon						
100 mM	CDS	1634 c.1634A>C GAG>GGC E545A <b>Transversion</b>	1658 c.1658delG AGT>ACC S553T Substitution	1659 c.1659T>C <b>Transition</b>			
300 mM	CDS	1570 c.1570A>T AGG>TGG R524W <b>Transversion</b>	1658 c.1658delG AGT>ACC S553T Substitution	1659 c.1659T>C <b>Transition</b>	3088 c.3088A>G ACT>GCT T1030A <b>Transition</b>	3226 c.3226A>G <b>Transition</b>	
500 mM	CDS	1623 c.1623T>C TCT>TCC S541S <b>Transition</b>	1634 c.1634A>C GAG>GGC E545A Substitution	1658 c.1658delG AGT>ACC S553T Substitution	1659 c.1659T>C <b>Transition</b>	3224 c.3224A>G <b>Transition</b>	

KRAS		Exon 1		Exon 2	
Control	CDS	89 c.89A>G GAC>GGC D30G <b>Transition</b>	144 c.144A>T GGA>GGT G48G <b>Transversion</b>	209 c.209A>G CAG>CGG E70R <b>Transition</b>	234 c.234T>C TTT>TTC F78F <b>Transition</b>
	Codon				
100 mM	CDS		183 c.183A>G CAA>CAG Q61Q <b>Transition</b>		
300 mM	CDS	86 c.86T>C GTG>GGC V29A <b>Transition</b>	188 c.188A>G GAG>GGG E63G <b>Transition</b>		
500 mM	CDS		162 c.162T>C GAT>GAC D54D <b>Transition</b>	285 c.285T>C CAT>CAC H95H <b>Transition</b>	

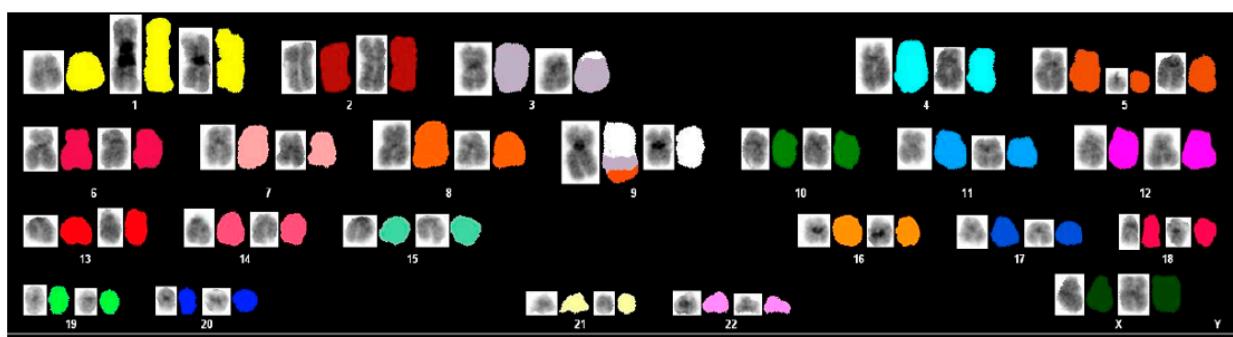
Transition	9
Transversion	4
Substitution	4

Transition	9
Transversion	1
Substitution	0

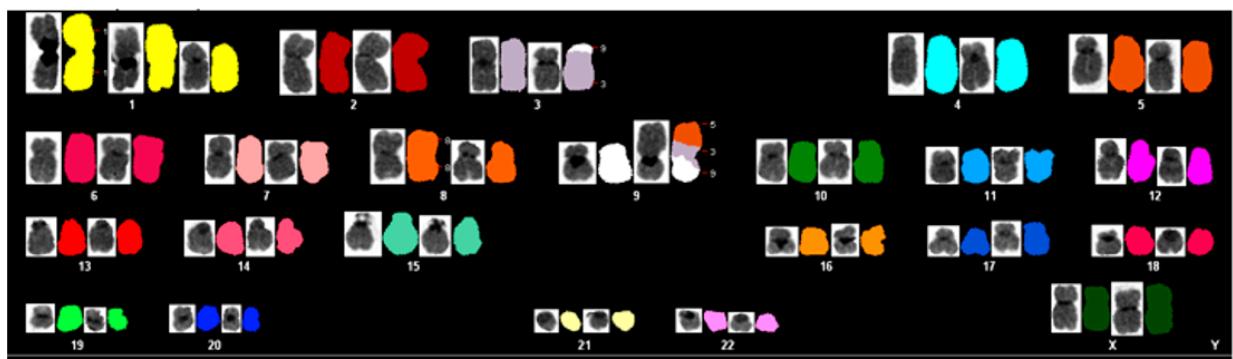
**Supplementary Figure S1:** Colony formation in SIT Media. Control and 100  $\mu$ M DETANO treated cells were evaluated for growth in serum-free media supplemented with 5 ng/ml selenium, 5 g/ml insulin, and 5 g/ml transferrin (SIT). 1x10<sup>5</sup> cells were plated in 60 mm dishes with SIT media and maintained for three weeks. When compared to control, MCF10A cells previously maintained in 100 $\mu$ M DETANO formed colonies in the SIT media.



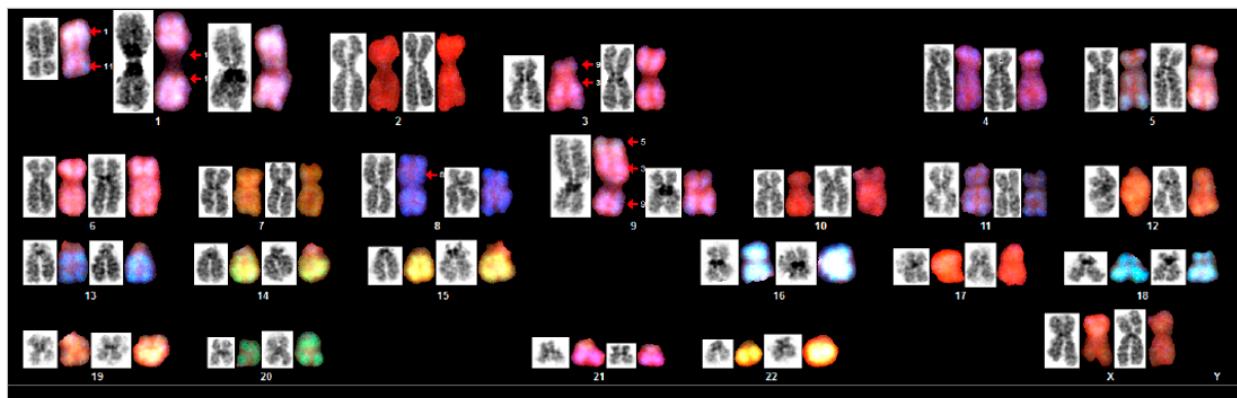
**Supplementary Figure S2: Cytogenetic Analyses.** Fig 4A. A dicentric Chr 1 composed by two Chr 1 long arm region 1, band 1; a derivative Chr 1; a translocation between Chr 3 short arm region 1, band 3 and Chr 9 short arm region 2, band 2; an isochromosome 8 form by two Chr 8 long arm region 1, band 1; a translocation involved partial Chr 9 Chr 3 & Chr 5; a deletion of Chr 21; and a translocation between Chr X short arm region 2, band 2 and Chr 21 long arm region 2, band 1. Fig 4B. Chromosome 21 was showed here in green by whole chromosome painting. The Chr 21 appeared normal in this FISH assay. TIMP1 is stained in orange. An Aqua X Control probe for region Xp13.2 stain. Fig 4C. Spectrum karyotyping of Control, 100 or 1000 mM DETANO treated MCF10A cells. Control: 47, XX, del (1), +i (1q), del (3), add (8), der (9) t (5; 3; 9). 100 uM: 47, XX, dic(1;1)(q11;q11), +der(1),t(3;9)(p13;p22), der(8) i(8)(q11;q11), t(5;3;9). 1000 uM: 47, XX, dic(1;1)(q11;q11), +der(1), t(3;9)(p13;p22), i(8)(q11;q11), t(3;5;9).



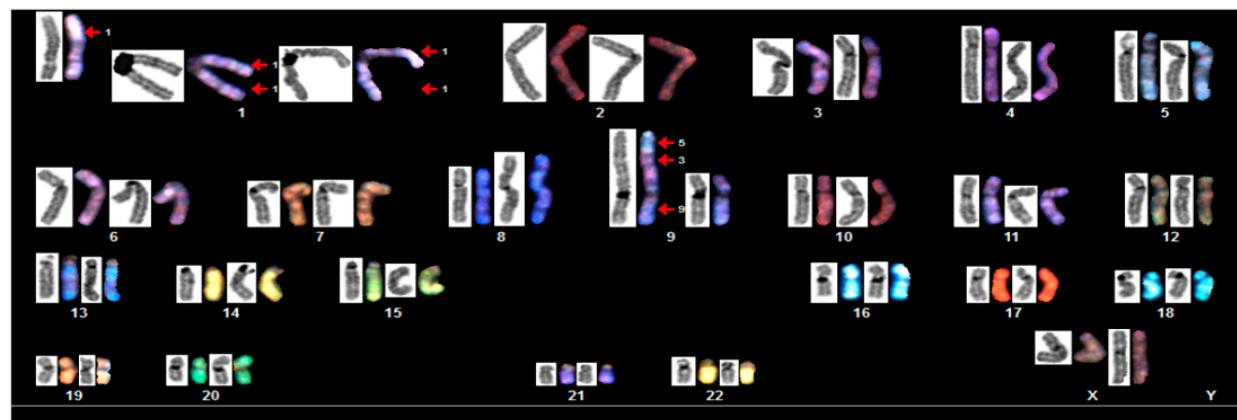
Control untreated; 47,XX, dic(1;1)(q11;q11),+der(1),-1,-3,t(3;9)(p13;22),-5,del(5),-8,i(8)(q11;q11),-9,t(9;3;5)



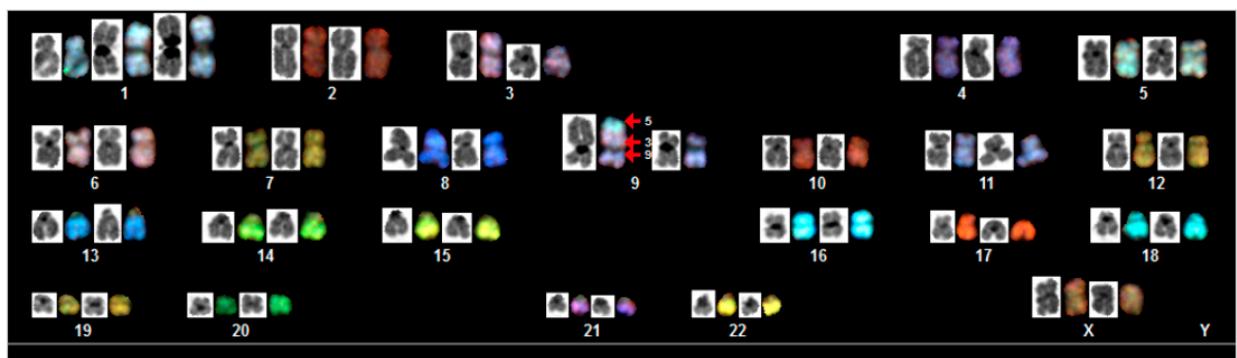
100 uM: 47, XX, dic(1;1)(q11;q11),+der(1),t(3;9)(p13;p22), der(8) i(8)(q11;q11), t(5;3;9)



2-1 100 uM: 47,XX, der(1) t(1;11),+dic(1q;1q),der(3)t(9;3),der(8),der(9)t(9;3:5)



2-2 100 uM: 47,XX, der(1),+(1q), der(3),der(8),t(9;3:5)



2-3 100 uM: 47,XX, der(1),+dic(1q;1q),der(3)t(9;3),der(8),der(9)t(9;3:5)