

ID	%GC	TM	Primer sequence	Best primer	Mapping(s)
LEFT_0	55.00	60.04	ATCTGACGCTGTTTGGGGAG		unique mapping
LEFT_1	55.00	60.11	CTAGAAGGTTTCCGCAGCGA		unique mapping
LEFT_2	60.00	60.32	GTC AAGCACCCACACCCTAG		unique mapping
LEFT_3	55.00	59.90	TCTTCCCTCTGCGCGTAATC		unique mapping
LEFT_4	60.00	60.46	GCCGAAACCTGATCCTCCAG		unique mapping
RIGHT_0	60.00	60.04	GACCTGCACTCCAATTCTCTC		unique mapping
RIGHT_1	50.00	60.04	AAGATCGTGCCAAGCGAAGA		unique mapping
RIGHT_2	60.00	60.74	CTGCAGTCCCCAAGATCGTG		unique mapping
RIGHT_3	57.90	59.85	TGGAAGGTGGCTGTGGTTTC		unique mapping
RIGHT_4	57.90	60.00	GTCCCCAAGATCGTGCCAA		unique mapping

55039274	CAGAACTTGGAGTCTGGCATCCACGCAGGGTGAGAGGCGGGAGAGGAGGCCCTAGGGCGCCGGCCTGCCTTCCAGC	
55039354	CCAGTTAGGATTTGGGAGTTTTTTTCTTCCCTCTGCGCGTAATCTGACGCTGTTTGGGAGGGCGAGGCCGAAACCTGATC	
	>>>>> LEFT_3 >>>>>	>>>>> LEFT_4
	>>>>> LEFT_0 >>>>>	
55039434	CTCCAGTCCGGGGTTCCGTTAATGTTTAATCAGATAGGATCGTCCGATGGGGCTCTGGTGCGTGATCTGCGCGCCCCA	
	>>>>>	
55039514	GGCGTCAAGCACCCACACCCTAGAAAGTTTCCGCAGCGACGTCGAGGCGCTCATGGTTGCAGGGGGCGCCGCCGTTTCA	
	>>>>> LEFT_2 >>>>>	
	>>>>> LEFT_1 >>>>>	
55039594	TTCAGGGTCTGAGCCTGGAGGAGTGAGCCAGGCAGTGAGACTGGCTCGGGCGGGCCGGGACGCGTCTGTCAGCAGCGGC	
55039674	TCCCAGCTCCCAGCCAGGATTCCGCGCGCCCTTCACGCGCCCTGCTCCTGAACCTTCAGCTCCTGCACAGTCTCCCCAC	

55039754	CGCAAGGCTCAAGGCGCCGC	CGGCGTGGACGCGCACGGCCTCTAGGTCTCCTCGCCAGGACAGCAACCTCTCCCCTGGC
	-----*	-----
55039834	CCTCATGGGACACGTGAGCTCCAGGCGGTCTGGTGCCGCTGCCACTGCTGCTGCTGCTGCTGCTGCTCCTGGGTC	
55039914	CGGGCGCCCGTGCGCAGGAGGACGAGGACGGCGACTACGAGGAGCTGGTGCTAGCCTTGCGTTCCGAGGAGGACGGCCTG	
55039994	GCCGAAGCACCCGAGCACGGAACACAGCCACCTTCCACCGCTGCGCCAAGGTGCGGGTGATAGGGATGGGAGGCCGGGGC	
	<<<<< RIGHT_3 <<<<<	
55040074	GAACCCGCAGCCGGGACGGTGCGGTGCTGTTTCTCTCGGGCCTCAGTTTCCCCCATGTAAGAGAGGAAGTGGAGTGCA	
	<<<<< RIGHT_0 <<	
55040154	GGTCGCCGAGGGCTCTTCGCTTGGCACGATCTTGGGACTGCAGGCAAGGCGGCGGGGAGGACGGGTAGTGGGAGCAC	
	<<<<	<<<<< RIGHT_1 <<<<<
		<<<<< RIGHT_4 <<<<<
		<<<<< RIGHT_2 <<<<<
55040234	GGTGGAGAGCGGGGACGGCCGGCTCTTTGGGGACTTGCTGG	

primer-designer version: 1.0-rc2 using dbSNP 150 for SNP checking, and human reference GRCh38.
Common SNP annotation: A common SNP is one that has at least one 1000Genomes population with a minor allele of frequency $\geq 1\%$ and for which 2 or more founders contribute to that minor allele frequency.