

Table S2 Primers and target sequences

Targets for disruption of genes

Gene Target	Name	Target Sequence
<i>Tet1</i>	gRNA Tet1Ex7	GGCATCGACCCAAAAACCTG CGG
<i>Tet2</i>	gRNA Tet2_9	GGAGCTGAGCCAAAAGAGGT TGG
<i>Tet3</i>	gRNA Tet3Ex7_2	GGCAAAGACCCTAACACCTG CGG

PCR primers for amplification of disrupted genes

Gene Target	Name	PCR Primer Sequence
<i>Tet1</i>	Tet1Ex7-1	CCATCTTCCAGGCGTACCT
	Tet1Ex7-2	AGGCACTGCAACAGGGTTAT
<i>Tet2</i>	Tet2Ex7-1	GCCAGAAGCAAGAAACCAAG
	Tet2Ex7-2	TGTTTAGGGGGTTAGCCAGA
<i>Tet3</i>	Tet3Ex7-3	TGCTTGGCCTATGTTATCCA
	Tet3Ex7-4	GCTCCGAGCATATTTGCAG

Targets for the deletion and inversion

Gene Target	Name	Target Sequence
<i>Tet1</i>	gRNA Tet1_2	GGCTGCTGTCAGGGAGCTCA TGG
	gRNA Tet1Ex7	GGCATCGACCCAAAAACCTG CGG

PCR primers for detection of deletion

Target	Name	Target Sequence
<i>Tet1</i> Ex4-Ex7 deletion	Tet1Ex7-2	AGGCACTGCAACAGGGTTAT
	Tet1Ex4-1	AGAACAAAGCCCCTGTGCTA

PCR primers for detection of inversion

Target	Primer	Target Sequence
<i>Tet1</i> Ex4-Ex7 inversion left	Tet1Ex7-1	CCATCTTCCAGGCGTACCT
	Tet1Ex4-1	AGAACAAAGCCCCTGTGCTA

Target	Primer	Target Sequence
<i>Tet1</i> Ex4-Ex7 inversion right	Tet1Ex7-2	AGGCACTGCAACAGGGTTAT
	Tet1Ex4-2	ACCACTCCAAGCCCTTTTCT

Target	Primer	Target Sequence
<i>Tet1</i> Control	Tet1RF-1	TTAGCTGGAGGGTGATCGTC
	Tet1RF-2	GTGGGCTGACGTGTGAAAA

*The PAM sequences are indicated in red.