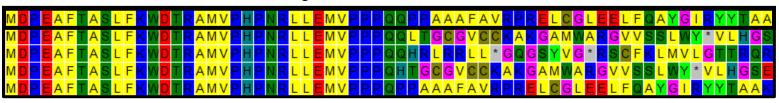
Amino acid modifications in PLFY for LFY-sg1



Amino acid modifications in PLFY for LFY-sg1sg2

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Genotype
WT
120 bp inversion
121 bp inversion
-120 bp
-121 bp
-130 bp

Amino acid modifications in PAG1 for AG-sg1sg2



Genotype
WT
+1 bp
-1 bp
-2 bp
-4 bp
-41 bp

WT
+1 bp
-1 bp
-2 bp
-3 bp

Amino acid modifications in PAG2 for AG-sg1sg2

MAYQNESQESSPL <mark>RK</mark> L	. GRGKVEIKRI	ENTINRQVIFCKRRN	G L L <mark>k k</mark> a <mark>y e</mark> l s v l c d a e v a l i v f s
MAYQNESQESSPLRKL	. <mark>GRGKVEIK</mark> AD) R E H H Q S S S H F L Q K A E	WFAQESL*IICSLRC*GCTHRLL
MAYQNESQESSPLRKL		SRTPPIV <mark>K</mark> SLSA <mark>KG</mark> GM	IVCS <mark>RKP</mark> MNYLFFAML <mark>R</mark> LHSSSSP
MAYQNESQESSPLRKL	. GRGKVESGSR	RTPPIAKSLSAKGGMV	'CS <mark>RKP</mark> MNYLFFAML <mark>R</mark> LHSSSS <mark>P</mark> A
MAYQNESQESSPLRKL	. G <mark>rgkveadr</mark> e	HHQS <mark>P</mark> SHFLQ <mark>K</mark> A E WF	AQESL*IICSLRC*GCTHRLLQP

Genotype
WT
+1 bp
-1 bp
-4 bp
-5 bp

Fig S1. Diversity in putative amino acid modifications to the WT peptide sequence in 717. These peptide alignments are the partial translation of the sequence alignment from the most common mutations seen in events with LFY-sg1, LFY-sg1sg2, and AG-sg1sg2. The first line in each alignment shows the WT sequence. Stop codons are shown with a *. The tables to the right identify the specific mutation that led to the peptide modification in each row.