



Allele Description

This is a CRISPR/Cas9 induced mutation creating a series of point mutations; V297M in exon ENSMUSE00001237825 of *HTRA1*. The stock was generated at MRC Harwell via microinjection of CRISPR/Cas9 reagents into 1-cell stage embryos.

qPCR Copy Counting Genotyping Strategy

The genotyping strategy presented here has been optimized for reagents and conditions used by the Genotyping Core at MRC Harwell. To genotype animals, we recommend researchers validate the assay independently. PCR cycling temperature and times may require additional optimization based on the specific genotyping reagents used.

Samples are genotyped using qPCR copy counting with both a wild type and a mutant assay against a known reference assay (*Dot1l* on chromosome 10; 2 copies present). Samples for this line are genotyped using the following primers and probe:

- Wild type (WT) assay with probe and reverse primer binding to the WT bases mutated in the mutant allele.
- Mutant assay with probe and reverse primer binding to the G601R, F606Y and R609H point mutations.

For autosomal genes that have been targeted, the following results would be expected:

Genotype of the Modified allele	WT Assay	Mutant Assay
Wildtype	2	0
Heterozygous	1	1
Homozygous mutant	0	2



HTRA1-V297M

HTRA1-V297M-WT1 assay (FAM labelled)

TCAGAGCTGAGACCTGGAGAATTTGTAGTTGCCATTGGAAGCCCCTTTTCTCTTCAAACACAG**GTCAC**
CACTGGGATCGTCAGCACCACCCAGCGAGGCG**GCAAAGAGCTGGGACTTCGGA**ACTCCGATATGG
ACTACATTCAGAC**CAGACGCTATCATCAATGTGAG**CCCTCTGCCCCCTCTACCTGCTGGGCTGGGGAC

Lower case letters denote bases changed in the mutant allele.
Probe sequence is in bold and shaded grey.
Primer sequences are in bold and underlined.

Oligo HTRA1-V297M	5' label	Sequence 5' → 3'	3' label	Oligo Type
HTRA1-V297M-WT_F	n/a	<u>GTCACCACTGGGATCGTC</u>	n/a	Wild type Forward
HTRA1-V297M-WT_PROBE	FAM	<u>AGTTCCGAAGTCCCAGCTCTTTGC</u>	ZEN/IBFQ	Wild type Probe
HTRA1-V297M-WT_R	n/a	<u>GATAGCGTCTGTCTGAATGTAGTC</u>	n/a	Wild type Reverse

HTRA1-V297M-MUT1 assay (FAM labelled)

TCAGAGCTGAGACCTGGAGAATTTGTAGTTGCCATTGGAAGCCCCTTTTCTCTTCAAACAC**CAGTCA**
CaACTGGGATCaTgAGCACCACCCAGCGAGGCG**GCAAAGAGCTGGGACTTCGGA**ACTCCGATATG
GACTACATTCAGAC**CAGACGCTATCATCAATGTGAG**CCCTCTGCCCCCTCTACCTGCTGGGCTGGGGGA

Lower case letters denote bases changed in the mutant allele.
Probe sequence is in bold and shaded grey.
Primer sequences are in bold and underlined.

Oligo HTRA1-V297M	5' label	Sequence 5' → 3'	3' label	Oligo Type
HTRA1-V297M-MUT_F	n/a	<u>CAGTCACA</u> ACTGGGATCATG	n/a	Mutant Forward
HTRA1-V297M-MUT_PROBE	FAM	<u>AGTTCCGAAGTCCCAGCTCTTTGC</u>	BHQ	Mutant Probe
HTRA1-V297M-MUT_R	n/a	<u>CTCACATTGATGATAGCGTCTG</u>	n/a	Mutant Reverse



Dot1l internal control (VIC labelled)

CTGATGGGTGTGGGCAGATCCTACAGAGTCCCATTGGCCACCATGTGTGCTACGCCTGAAATAAAGCCTT**GCC**
CCAGCACGACCATTCAGGG**CCAGCTCTCAAGTCG**ACTGTAAGATGAAGCATAAGGATGCCAACTACTAACA
GAAAACGACTAGAGGGGAAAAGAACAAGGAAACAGAAGACGCAGCACTCCGGCTTCCCTGGGTTGGCCAGT
CACCTATGA

Oligo HTRA1-V297M	5' label	Sequence 5' → 3'	3' label	Oligo Type
Dot1l_Forward	n/a	<u>GCCCCAGCACGACCATT</u>	n/a	WT Forward
Dot1l_Probe	VIC	CCAGCTCTCAAGTCG	BHQ	WT Probe
Dot1l_Reverse	n/a	<u>TAGTTGGCATCCTTATGCTTCATC</u>	n/a	WT Reverse

Probe sequence is in bold and shaded grey
Primer sequences are in bold and underlined

DNA extraction method

DNA is extracted from ear clips using Applied Biosystems Taqman Sample-to-SNP Kit and qPCR run using 1:10 dilution from the crude preparation.

qPCR master mix 1X

Applied Biosystems GTX Taqman master mix	5 µl
Dot1l_Forward (20 µM)	0.225 µl
Dot1l_Reverse (20 µM)	0.225 µl
Dot1l_Probe (5 µM)	0.2 µl
FAM Assay (probe 5 µM & primers 15 µM each)	0.3 µl
ddH2O	1.55 µl
DNA (1:10 dilution of ABI Sample-to-SNP prep)	2.5 µl

Each sample is ran in technical duplicate. Seven WT and/or mutant controls are also included in duplicate along with non-template controls.

qPCR cycling conditions

qPCR instrument: Applied Biosystems 7500/7900 or ThermoFisher QuantStudio 7

95°C for 20 sec
Then 40 cycles of;
95°C for 3 sec
60°C for 30 sec

