

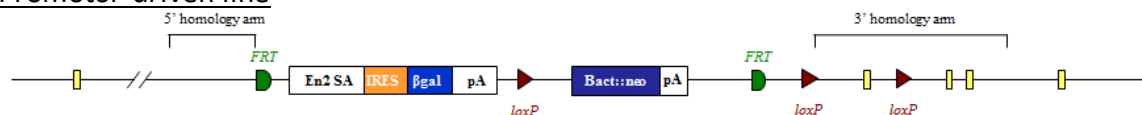


## EUCOMM/KOMP-CSD 'Knockout-First' Genotyping

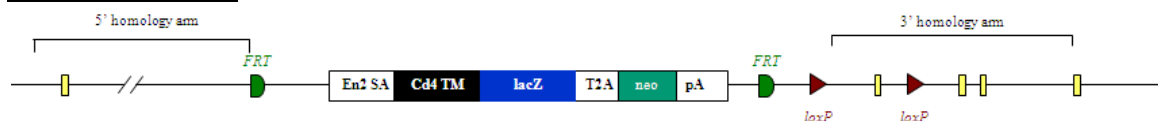
### Introduction

The majority of animals produced from the EUCOMM/KOMP-CSD ES cell resource contain the Knockout-First-Reporter Tagged Insertion allele. As well as gene specific elements, the targeting cassette contains many common elements e.g. encoding beta-galactosidase (lacZ gene), neomycin-resistance (neo gene), FRT and loxP sites. Further details can be found on <http://www.knockoutmouse.org/about/eucomm>. The majority of cassettes will be either promoter driven where the neo gene contains its own promoter and is separated from the lacZ gene by a 3<sup>rd</sup> loxP site, or promoterless (PL) where the neo and lacZ genes are adjacent and there are only 2 loxP sites. These have further implications in our genotyping strategies detailed later.

### Promotor driven line



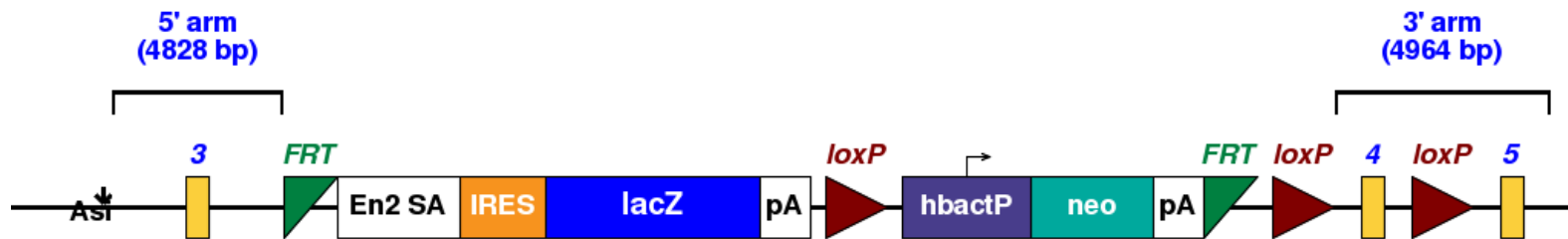
### Promotorless line



Animals containing the full Knockout-First mutant allele (Tm1a) above can be crossed to mice expressing Flp or Cre recombinase. The progeny from these mating may contain converted forms of the Tm1a allele known as Tm1b, Tm1c and Tm1d depending on the breeding strategy performed. In order to genotype animals generated from the EUCOMM/KOMP-CSD 'Knockout-First' ES cell resource, a combination of mutant assays looking for sequences occurring in the targeting cassette, and WT specific assays will need to be performed. We typically use real time qPCR and copy count the various alleles. This assay design sheet will give an outline of how to genotype these lines by gel based and qPCR based technologies, as well as including details of assays we have tried and tested.



## Pla2g2c-Tm1a



Details for the tm1a allele can be found below

<https://www.mousephenotype.org/data/genes/MGI:106638>



## Types of alleles

### Promotor driven lines

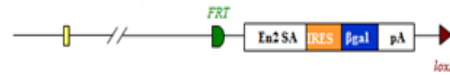
**WT** allele



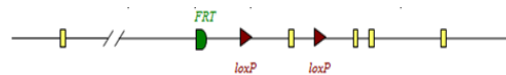
**Tm1a** promotor driven construct



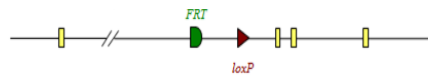
**Tm1b** created by action of Cre recombinase on Tm1a allele



**Tm1c** created by action of Flp recombinase on Tm1a allele



**Tm1d** created by action of Cre recombinase on Tm1c allele



### Promotorless lines

**WT** allele



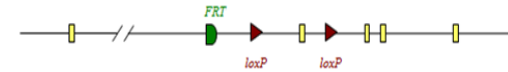
**PL-Tm1a** (promotorless) construct



**PL-Tm1b** created by action of Cre recombinase on PL-Tm1a allele



**Tm1c** created by action of Flp recombinase on PL-Tm1a allele



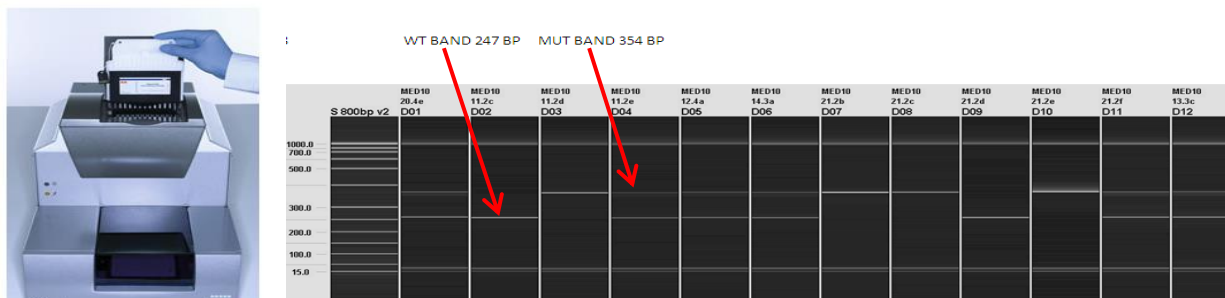
**Tm1d** created by action of Cre recombinase on Tm1c allele





## Gel based genotyping

The gel based assays are normally run on the Qiagen QIAxcel. This is a capillary based system that provides clearer resolution and is quicker than running standard agarose gels. Different size ladders maybe loaded onto runs depending on the fragment sizes being analysed. Typically samples are run with a 50-800bp size ladder.



PCR is performed using KAPA2G Fast Hotstart Readymix (2X), although alternatives may be used.

### Generic 4 primer PCR mix

KAPA2G Fast Hotstart Readymix (2X)	5µl
Primer 1F (20µM)	0.5µl
Primer 1R (20µM)	0.5µl
Primer 2F (20µM)	0.5µl
Primer 2R (20µM)	0.5µl
H <sub>2</sub> O	2.0µl
DNA (~30ng)	1µl

### 60°C Cycling conditions

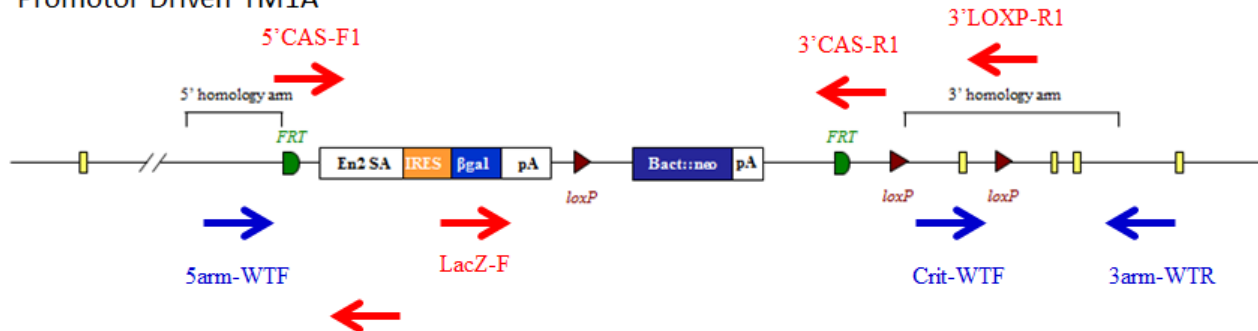
<b>60TM30FA</b>		
1.	95°C	1min
2.	95°C	10sec
3.	<b>60°C</b>	10sec
4.	72°C	1sec
5.	Go to 2 for 29 cycles	
6.	72°C	30sec
7.	16 °C	forever



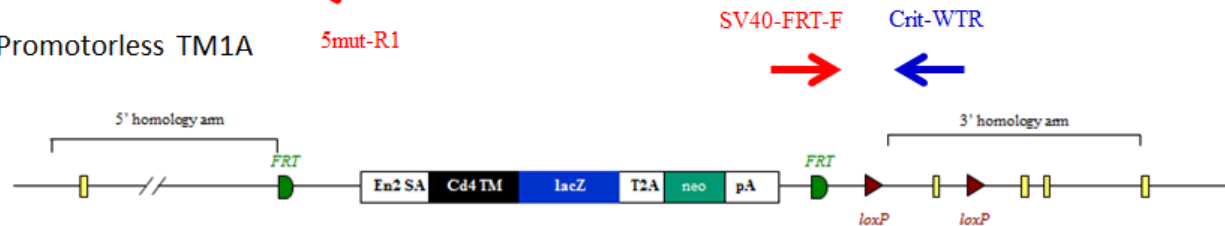
## Types of gel based assay

The diagram below shows the location of the various gel based primers.

### Promotor Driven TM1A



### Promotorless TM1A



## Mutant primer sequences

5mut-R1	GAACTTCGGAATAGGAACTTCG
LacZ-F	CCAGTTGGTCTGGTGTCA
SV40-FRT-F	CGCGTCGAGAAGTTCCTATT
5'CAS-F1	AAGGCGCATAACGATACCAC
3'CAS-R1	CCGCCTACTGCGACTATAGAGA
3'LOXP-R1	ACTGATGGCGAGCTCAGACC



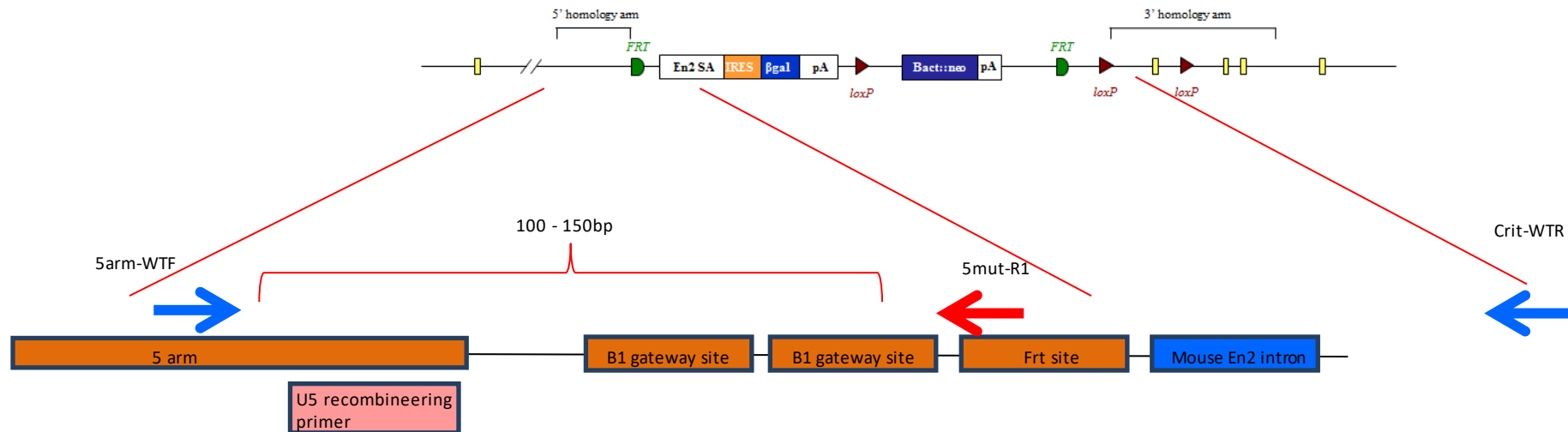
## Tm1a gel based genotyping

Tm1a gel based assays are designed using a universal mutant reverse primer that sits in the sequence just after the 5' homology arm

5mut-R1                    GAACTTCGGAATAGGAACTTCG

A forward primer is designed to the 5' homology arm and designed to give a mutant specific band, typically between 100-150bp that will only be present if the cassette is present.

A WT reverse primer is designed to the critical region and should give a product of >200bp. If the mutant cassette is present the product between the two primers is too large to PCR under standard conditions.





## Pla2g2c Tm1a gel based genotyping

### Pla2g2c 'Knockout'-First and WT sequences

#### 5'homology arm (last 300bp)

TTCTCCTGTGGGTGGACATATGACATTTCCAGTGTCTGATCTTACCACGCTGCCAGCAAACACCCTTGTATATGCATCTCTGCGTTGTTGTGTGAGCATTTTGTAAAAGTGAAA  
AGTCTACTGATGCATGCCTGAGAAGTCCAGCACCTCGAAGGTTTGGACTGGTCTCTATTGGGTAAGACAAGACCCTGTCCGTTAAAACAAAATAAGAATTGGGTCTGTGTGATT  
TGCCATTTTATAGGCTGGATCAAGCGTGTCTTACTTTCTTACCCTGTGTTCTGCTCCTGTGAGAC

#### Mutant 5'sequence location of 5mut-R1 (mutant reverse primer)

AAGGCGCATAACGATACCACGATATCAACAAGTTTGTACAAAAAAGCAGGCTGGCGCCGGAAC **CGAAGTTCCTATTCCGAAGTTC** CTATTCTCTAGAAAGTATAGGAACTTCGAAC  
CCTTTCCCACACCACCCTCCACACTTGCCCCAAACACTGCCAAC

#### Critical region

CAGCCATGCGACCCGTTGACCTTGTGAGCAGACGATGCACCTTCTTCTTAAATGCTACCTGTACTCTCAGTCTAGAGGGCCTAACAGGATCAGTGTCAACAGAGAAAACACCGGCT  
ACAGACTGGAACCTGTGGCTTTGAATTTTCCAGTTAACGTTGTTTCCAGACTGGATGGTGGTAGAGCAACCCCTGACAGCTGGGGAGCCTGAGCCCCATCCTGATTAGCAAGACAC  
TTTCTGAGTGTGGAGTAACACCAGTCTAAATTTCTTCTTGAAGGAGGCTGATGTCTTGGGGGAGGGTCTGGGCTGTCCGCCCTCTGCAAACACTCCAGCAAAGGCCGTGG  
GCCCTCACACTCTTCTTGTGTTTATTGTTTCTCCTTGGCAGGGACAACCTCCACCCTCAGCAGTTTCTGGCAGTTCAGAGGATGGTCAAACACGTCACAGGGCGCAGCGCCTTCTT  
CTCCTATTACGGATATGGCTGCTACTGTGGGCTTGGGGGCAAAGGGCTCCCTGTGGATGCCACAGACAGGTGATGCGGATGACCTGGGCCAACCTTATTTTATTGCTTTGAAGT  
AGAGGCTTTACAGCCAGGGGAGAGGGTGTGGGAATCCTGAAGAAACAATTACAGGACAGCACAAAGATGAAGAAGAGTTGCAGGGTCAGTCCCGAGGAAGCCAGAGCTGAG

#### 3'homology arm (1st 300bp)

TGAGACAGGGTCTTATCTTCCAACCTCACGTTGGCTCCTAACTTACTATATAGTCCAGGTTGGACTTGAACCTCTGGACCCTCCTGCCTCAGCCACCCAAGTGTTTCCAGATTCAGGCA  
TATGCCACTAGGTACCGCTCAAACCCAGAACCTCATCCCTGCTGGACAGAGATTACCCCAAGCCCCATCCTGCTAGCCTCCATCCAGGACCCAGTGTTACACAGTCAGTCTTGCC  
CAGAGCTCTGGAGGCCTCACCTATTTGCTTACAAAGGTGATCTTTCTCATCTGCTCCATTTTTTTTTT



## Pla2g2c Tm1a gel based genotyping

### Pla2g2c 'Knockout'-First and WT sequences

#### Genomic sequence

Black sequence is missing in the construct due to the insertion of loxP sites and can be used to design break point (BP) qPCR assays

```
TTCTCCTGTGGGTGGACATATGACATTTCCAGTGTTCTGATCTTACCACGCTGCCAGCAAACACCCTTGTATATGCATCTCTGCGTTGTTGTGTGAGCATTTTGTAAAAGTGAAAA
AGTCTACTGATGCATGCCTGAGAAGTCCAGCACCTCGAAGGTTTGGAGACTGGTCTCTATTGGGTAAGACAAGACCCTGTCCGTTAAAACAAAATAAGAATTGGGTCTGTGTGATT
TGCCATTTTATAGGCTGGATCAAGCGTGTTTTACTTTCTCTTACCCCTGTGTTCTGCTCCTGTGAGACTCTAACGGCCTGATGCTACCTAGAGGAGGGACTGACTGGAACATCAGC
CTCTCTGTTGAGGGTCTCTCAGGTAGCTGACTTCTAAGGAGCTGTGCACAGCCATGCGACCCGTTGACCTTGTGAGCAGACGATGCACCTTCTTCTTAAATGCTACCTGTACTCTC
AGTCTAGAGGGCCTAACAGGATCAGTGTCAACAGAGAAAACACCCGGCTACAGACTGGAACCTGTGGCTTTGAATTTCCAGTTAACGTTGTTTCAGACCTGGATGGTGGTAGAGCA
ACCCCTGACAGCTGGGGAGCCTGAGCCCCATCCTGATTAGCAAGACACTTCTGAGTGTGGAGTAACACCAGTCTAAATTTCTTTCCTTTGAAGGAGGCTGATGTCTTGGGGGAG
GGTTCTGGGCTGTCCGCCCTCTGCAAACACTCCAGCAAAGGCCGTGGGCCCTCACACTCTTCTTGTATTATTGTTTCCCTTGGCAGGGACAACCTCCACCCTCAGCAGTTTCTG
GCAGTTCAGAGGATGGTCAAACACGTACAGGGCGCAGCGCCTTCTTCTCCTATTACGGATATGGCTGCTACTGTGGCTTGGGGGCAAAGGGCTCCCTGTGGATGCCACAGACA
GGTGATGCGGATGACCTGGGCCAACCCCTATTTTTATTTGCTTTGAAGTAGAGGCTTTACAGCCAGGGGAGAGGGTGTGGGAATCCTGAAGAAACAATTACAGGACAGCACAAGAT
GAAGAAGAGTTGCAGGGTCAGTCCCGAGGAAGCCAGAGCTGAGCCACCTTCCCTCTGTGTGACCTGTGACACTGCCTAGGGGCCGTGATGCCTGGGATCCTTAGTATGGCTTGTGTG
CTGTTGCCATCTTGAGACAGGGTCTTATCTTCCAACCTCACGTTGGCTCCTAACTTACTATATAGTCCAGGTTGGACTTGAACCTCTGGACCCTCCTGCCTCAGCCACCCAAGTGTTC
AGATTCCAGGCATATGCCACTAGGTACCGCTCAAACCCAGAACCTCATCCCTGCTGGACAGAGATTACCCCAAGCCCCATCCTGCTAGCCTCCATCCAGGACCCCAAGTGTACACA
GTCAGTCTGCCAGAGCTCTGGAGGCCTCACCTATTTGCTTACAAAGGTGATCTTTCCTCATCTGCTCCATTTTTTTTTT
```

#### 5' homology arm and cassette sequence

```
TTCTCCTGTGGGTGGACATATGACATTTCCAGTGTTCTGATCTTACCACGCTGCCAGCAAACACCCTTGTATATGCATCTCTGCGTTGTTGTGTGAGCATTTTGTAAAAGTGAAAA
AGTCTACTGATGCATGCCTGAGAAGTCCAGCACCTCGAAGGTTTGGAGACTGGTCTCTATTGGGTAAGACAAGACCCTGTCCGTTAAAACAAAATAAGAATTGGGTCTGTGTGATT
TGCCATTTTATAGGCTGGATCAAGCGTGTTTTACTTTCTCTTACCCCTGTGTTCTGCTCCTGTGAGACAAGGCGCATAACGATACCACGATATCAACAAGTTTGTACAAAAAAGCA
GGCTGGCGCCGGAACCGAAGTTCCATTTCCGAAGTTCCATTTCTCTAGAAAGTATAGGAACTTCAACCCCTTTCCACACCACCCTCCACACTTGCCCCAAACACTGCCAAC
```





## Tm1a gel based genotyping

### Pla2g2c tm1a gel based primers

Pla2g2c-5arm-WTF TGATGCATGCCTGAGAAGTC  
Pla2g2c-Crit-WTR ACCCTCAACAGAGAGGCTGA

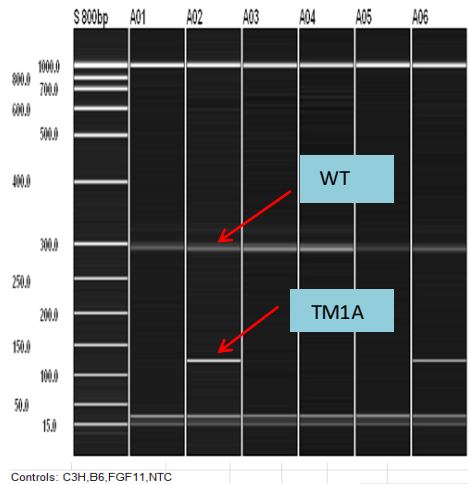
CAS\_R1\_Term TCGTGGTATCGTTATGCGCC

WT band 240  
Mutant band 199

### PCR mix

KAPA2G Fast Hotstart Readymix (2X) 5µl  
5arm-WTF (20µM) 0.5µl  
Crit-WTR (20µM) 0.5µl  
CAS\_R1\_Term (20µM) 0.5µl  
H<sub>2</sub>O 2.5µl  
DNA (~30ng) 1µl

### Example of a generic TM1A assay



### Cycling conditions

#### **60TM30FA**

1. 95°C 1min.
2. 95°C 10sec.
3. 60°C 10sec.
4. 72°C 1sec.
5. Go to 2 for 29 cycles
6. 72°C 30sec.
7. 16 °C forever
8. end

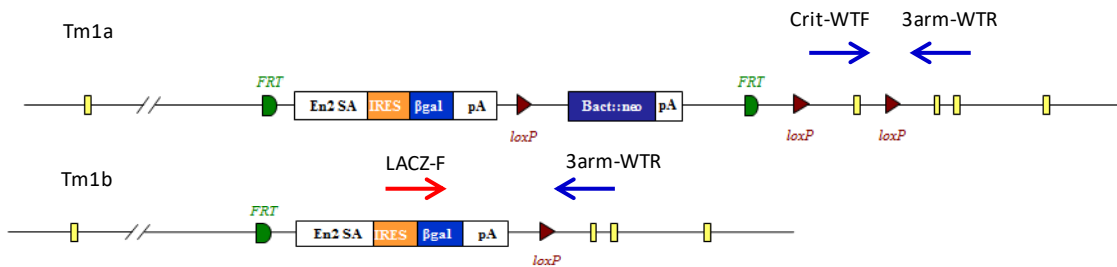


# Pla2g2c

## Tm1b gel based genotyping

### Promotor driven genotyping

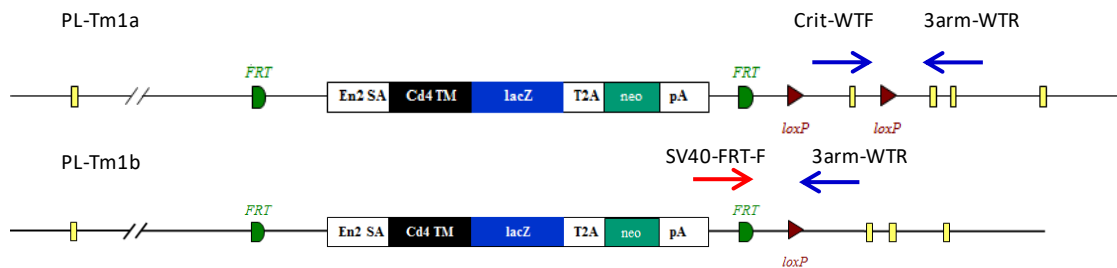
Tm1b alleles are produced by exposing tm1a alleles to cre recombinase which causes the deletion of Neo and the critical region.



If the deletion of the critical region has not occurred then a product is produced between the -Crit-WTF and the -3arm-WTR. Complete conversion by Cre will mean a product is produced from the LacZ-F and -3arm-WTR as the two primers now lie close enough to generate a PCR product under standard conditions. Run both the tm1a genotyping protocol to detect WT and tm1a alleles and the tm1b protocol to detect converted alleles.

### Promotorless genotyping

PL-Tm1b (promotorless tm1b) mice are produced in a similar fashion but only the deletion of the critical region occurs and Neo is still present.



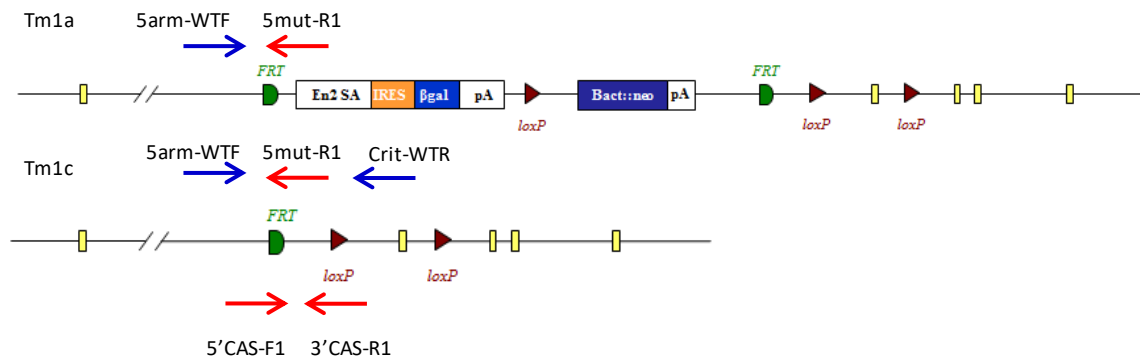
If the deletion of the critical region has not occurred then a product is produced between the Crit-WTF and the 3arm-WTR. Complete conversion by Cre will mean a product is produced from the SV40-FRT-F and 3arm-WTR as the two primers now lie close enough to generate a PCR product under standard conditions. This will not work for promotor driven lines as the presence of the 5' loxP site between the Neo and lacZ sequence causes this region to be removed by cre deletion. Run both the tm1a genotyping protocol to detect WT and tm1a alleles and the tm1b protocol to detect converted alleles.



## Tm1c and Tm1d gel based genotyping

### Tm1c genotyping

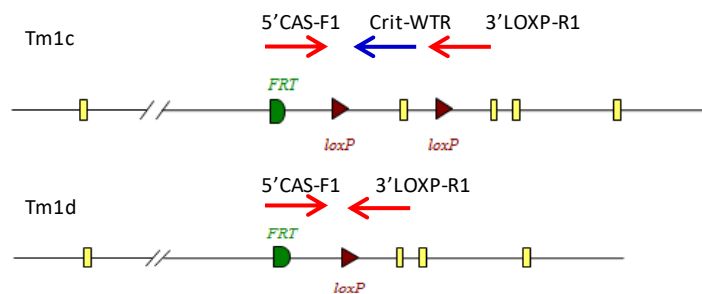
Tm1c alleles are produced by exposing tm1a alleles to flp recombinase which causes the deletion of Neo and LacZ sequences that are between the FRT sites.



The protocol for tm1a genotyping needs to be run. Both tm1a and tm1c alleles will produce a mutant specific product from -5arm-WTF and 5mut-R1. WT and tm1c alleles will produce PCR products from -5arm-WTF and Crit-WTR although the tm1c product will be around 200bp bigger due to the presence of the FRT-loxP part of the cassette. In addition to this a generic tm1c assay can be used. This uses the 5' CAS-F1 and 3' CAS-R1 primers run with an internal control. The tm1c allele produces a 218bp product.

### Tm1d genotyping

Tm1d mice are produced by crossing tm1c animals with mice expressing cre recombinase. This causes the deletion of the critical region.



The tm1d allele produces a 174bp product from the 5' CAS-F1 and 3' LOXP-R1 primers. A tm1c allele will produce a larger product which includes the critical region sequence. If the critical region is very large then a gene specific primer lying in the critical region may be needed to detect it (Crit-WTR).



## Gel based genotyping strategy summary

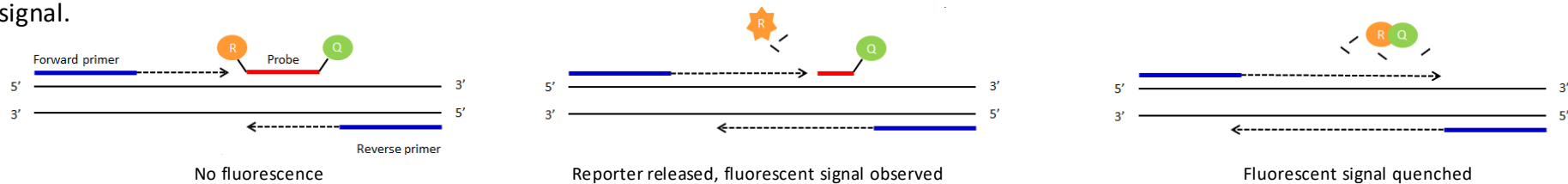
	Primers to use										Comments
	5'CAS-F1	5mut-R1	LacZ-F	SV40-FRT-F	3'CAS-R1	3'LOXP-R1	5arm-WTF	Crit-WTF	Crit-WTR	3arm-WTR	
<b>Tm1a + PL</b>		✓					✓		✓		
<b>Tm1b</b>			✓					✓		✓	Run tm1a protocol to identify WT alleles
<b>PL-Tm1b</b>				✓				✓		✓	Run tm1a protocol to identify WT alleles
<b>Tm1c</b>	✓	✓			✓		✓		✓		Blue =separate reaction
<b>Tm1d</b>	✓					✓			✓		Run tm1a protocol to identify WT alleles

- WT Product between 5arm-WTF and Crit-WTR (product too big in tm1a).
- Tm1a Product between 5arm-WTF and 5mut-R1.
- Tm1b Product between LacZ-F and 3arm-WTR (product too big in tm1a but PCR product from Crit-WTF and 3arm-WTR is made).
- Tm1b-PL Product between SV40-FRT-F and 3arm-WTR (product too big in tm1a but PCR product from Crit-WTF and 3arm-WTR is made).
- Tm1c 218bp product between 5'CAS-F1 and 3'CAS-R1 made. Can also run tm1a protocol for a gene specific assay.
- Tm1d 174bp product produced from 5'CAS-F1 and 3'LOXP-R1. Larger product may be produced by tm1c.



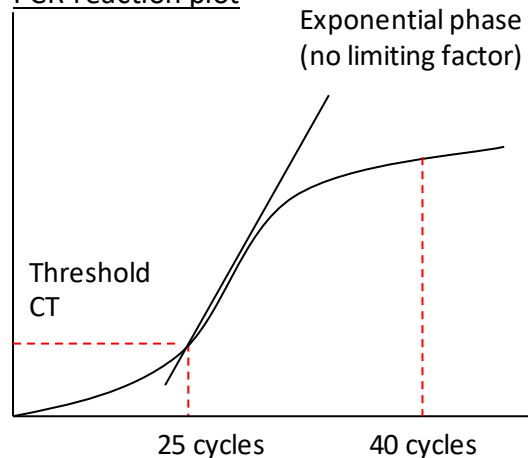
## qPCR based genotyping

Standard PCR is the amplification of DNA between a pair of primers. Quantitative PCR employs the same principal as standard PCR, although it actually monitors the progress of the DNA synthesis as it occurs. The progress of the reaction is measured by using a Taqman probe. This is a short DNA oligo that is complimentary to part of the DNA sequence between the forward and reverse primers. At the 5' end of the probe there is a fluorescent reporter (R) and at the 3' end a quencher (Q). Whilst they are in close contact with each other there is no fluorescent signal.



As the forward primer is extended the reporter is cleaved from the probe resulting in a fluorescent signal being detected. Once the primer extends enough to release the quencher this signal is blocked. By using probes with different fluorescent signals multiple PCR assays can be multiplexed and run together.

### PCR reaction plot



The number of cycles the PCR takes to reach a set threshold is known as the CT value. This is inversely correlated to the amount of template DNA in the sample.

e.g. CT 25 = 2 x template DNA

CT 26 = 1 x template DNA

CT above 30 = no template represented in the sample

CT value can be used to determine how many copies of a particular allele samples have.



## qPCR assay types

All qPCR assays are FAM labelled and run in duplex with a VIC labelled internal control, Dot1l. There are 4 mutant qPCR assays based around the targeting cassette. They are as follows:

- : Neomycin assay                      Designed around the sequence of the neomycin resistance cassette.
- : LacZ assay                              Designed around the sequence of the LacZ reporter.
- : Promotor driven Tm1b                Designed to detect the recombined sequence when a promotor driven tm1a allele is converted to a tm1b allele. The forward primer is based on the region of sequence of the cassette between LacZ and Neo sequences just upstream of the 5' most loxP. This is not present in promotorless alleles. The reverse primer is based on sequence found in the 3' loxP region.
- : FRT-loxP                                Designed to detect the recombined sequence of the FRT and loxP area created when the Tm1a allele is Flp converted to the tm1c. The forward primer is based on the sequence of the 5' region upstream of the 5' FRT site which means this assay will only work for Tm1c/d.

There are 2 WT qPCR assays based on WT sequence that may be lost in Tm1a/b/c/d alleles. They are as follows:

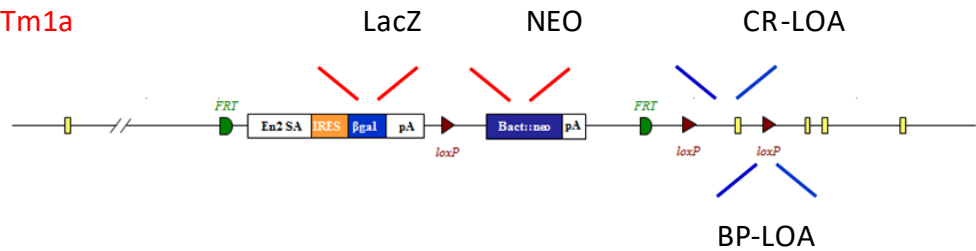
- : BP-LOA                                  Break Point loss of allele assay. This is based on WT sequence that is lost around the loxP sites of the synthetic cassette. This detects any cassette and should drop in copy number if a tm1a/b/c or d allele is present.
- : CR-LOA                                  Critical Region loss of allele assay. This is based on the critical region sequence that is flanked by the two loxP sites. This will only drop in copy number if a tm1b or tm1d is present.



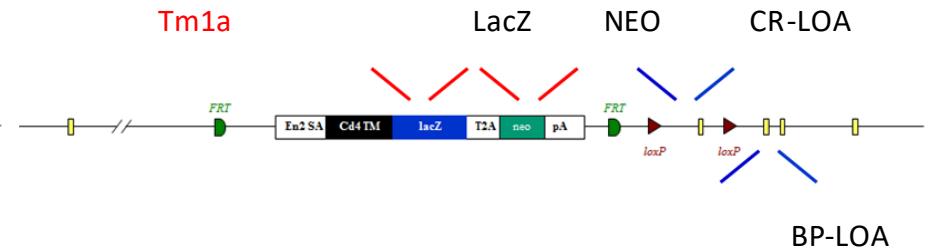
# Pla2g2c

## qPCR based methods to distinguish alleles

Promotor Driven  
Tm1a

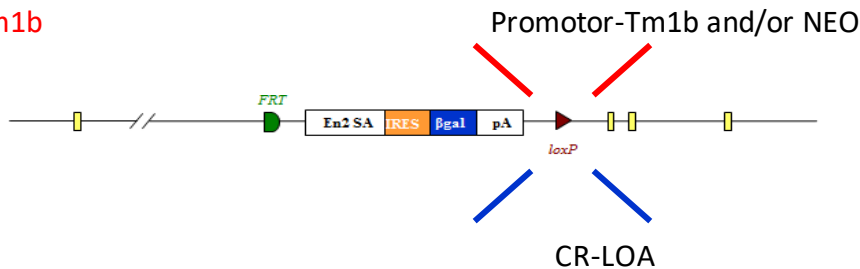


Promotorless  
Tm1a

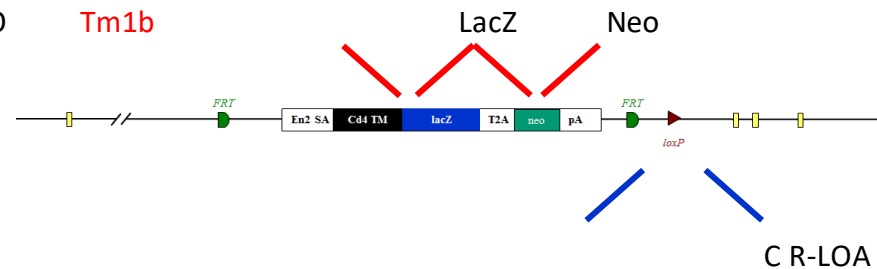


Run either LacZ or Neo which will identify the cassette and the BP-LOA which will identify the correct line as this should drop in copy number if the cassette has replaced the WT allele. The critical region should be 2 copies and may be run to distinguish tm1a alleles from tm1b alleles. Both the promotor driven tm1b and FRT-loxP should not work as they detect recombined cassettes (tm1 b/c/d) not the complete cassette (tm1a).

Promotor Driven  
Tm1b



Promotorless  
Tm1b

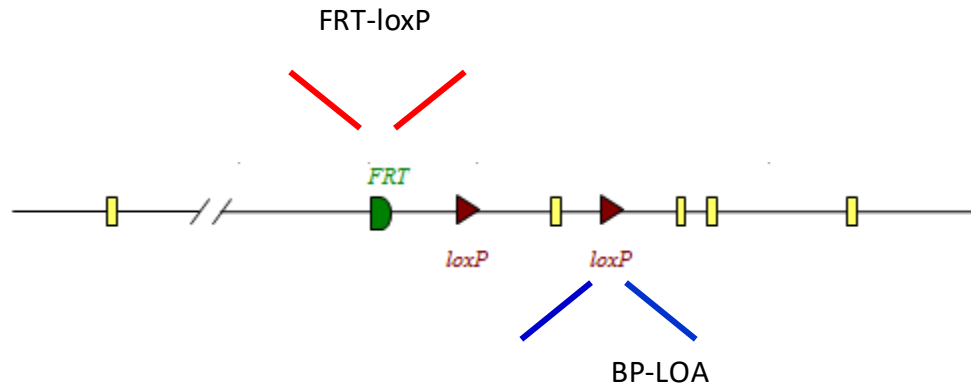


For both promotor driven and promotorless lines a drop in the CR-LOA copy number identifies the correct gene has been targeted. For promotor driven lines if the CR drops and the promotor-tm1b assay calls positive by the same value as the drop then the allele has been fully converted to a tm1b. A drop in Neo and the CR-LOA also would signify the same thing. For promotorless lines just a drop in the CR is enough to identify a tm1b or d allele and the presence of Neo or LacZ indicates it is not a tm1d.



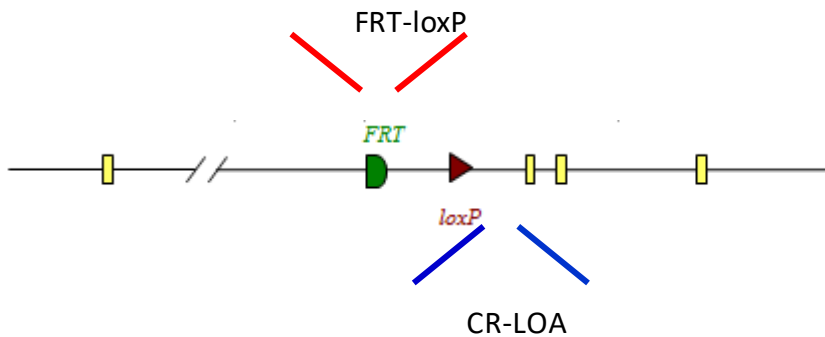
## qPCR based methods to distinguish alleles continued

### Tm1c



A positive result for FRT-loxP confirms a tm1c or d allele. Running the BP-LOA and a drop in copy number will confirm the correct line has been targeted. To distinguish between tm1c and a tm1d allele (below) the CR-LOA must also be run. A drop in copy number for the CR-LOA indicates a tm1d allele.

### Tm1d







## qPCR mutant assay sequences

### NEO assay (FAM labelled probe)

ATTGAACAAGATGGATTGCACGCAGGTTCTCCGGCCGCTTG **GGTGGAGAGGCTATTCGGC** TATGAC **TGGGCACAACAGACAATCGGCTC** CT **CTGATGCCGCCGTGTTCC** CGGCTGTC  
AGCGCAGGGGGCGCCCGTTCTTTTTGTCAAGACCGACCTGTCCGGTGCCCTGAATGAACTGCAGGACGAGGCAGCGCGGCTATCGTGGCTGGCCACGACGGGCGTTCCTTGCGCAG  
CTGTGCTCGACGTTGTCACTGAAGCGGGAAGGACTGGCTGCTATTGGGCGAAGTGCCGGGGCAGGATCTCCTGTACATCTCACCTTGCTCCTGCCGAGAAAGTATCCATCATGGCT  
Primer 1 = GGTGGAGAGGCTATTCGGC      Primer 2 = GAACACGGCGGCATCAG      Probe = TGGGCACAACAGACAATCGGCTG

### LacZ assay (FAM labelled probe)

CGATAAAAAACAACCTGCTGACGCCGCTGCGCGATCAGTTACCCGCTGCACCGCTGGATAACGACATTGGCGTAAGTGAAGCGACCCGATTGACCCTAACGCCTGGGTGCGAACGCT  
GGAAGGCGGGCGGCCATTACCAGGCCGAAGCAGCGTTGTTGCAGTGCACGGCAGATACTTGTGATGCGGTGCTGATTACGACCGCTCACGCGTGGCAGCATCAGGGGAAAACC  
TTAT **TTATCAGCCGGAAAACCTACC** **GGATTGATGGTAGTGGTCAAATGGCGA** TTACC **GTTGATGTTGAAGTGGCGAG** CGATACCCGCATCCGGCGCGGATTTGGCCTGAACTGCCA  
GCTGGCGCAGGTAGCAGAGCGGGTAAACTGGCTCGGATTAGGGCCGCAAGAAAATATCCCGACCGCCTTACTGCCGCCTGTTTTGACCGCTGGGATCTGCCATTGTCAGACATGT  
Primer 1 = CTCGCCACTTCAACATCAAC      Primer 2 = TTATCAGCCGGAAAACCTACC      Probe = TCGCCATTTGACCACTACCATCAATCC

### Promotor-Tm1b-Mut1 (FAM labelled probe)

ATTGTTGTTGTTAACTTGTTTATTGCAGCTTATAATGGTTACAAATAAAGCAATAGCATCACAAATTCACAAATAAAGCATTTTTTT **CACTGCATTCTAGTTGTGGTTTTGTC** CAA  
ACTCATCAATGTATCTTATCATG **TCTGGATCCGGAATAACTTCGTATAGCA** TACATTATACGAAGTT **ATGGTCTGAGCTCGCCATCAGT** TCA  
Primer 1 = CACTGCATTCTAGTTGTGGTTTTGTC      Primer 2 = ACTGATGGCGAGCTCAGACCAT      Probe = TCTGGATCCGGAATAACTTCGTATAGCA

### FRT-LOXP-mut2 assay (FAM labelled probe)

**AGGCGCATAACGATACCACGATA** TCAACAAGTTTGTACAAAAAGCAGGCTGGCGCCGGAACCGAAGTTCCTAT **TCCGAAGTTCCTATTCTCTAGA** AAGTATAGGAAC **TTC<GTC**  
**GAGATAACTT>CGTATAGCA** TACATTATACGAAG  
Primer 1 = AGGCGCATAACGATACCACGATA      Primer 2 = TGCTATACGAAGTTATCTCGACGA      Probe = TCCGAAGTTCCTATTCTCTAGA



## qPCR WT assay sequences

### Example of a BP-LOA-wt1 assay (FAM labelled probe)

Blue = Sequence in homology arm    Black = breakpoint    Red = critical region

TACAACCTCTGTATGAGCCCATTCCTCCTCCTGGAAGAACACATATTTGGATTGATTTTAACTCGGATTTTGCAGGCTGCTTTTGCACCTGAAACGTTTTTAAATATATTAC<CCAA  
ACCAATCTTGGCAAGTTAGCAAGCCTTTTTAGAGACACATCGGTGAGTCGGTGGTTTGTGAAACACTC>CGTCATTCTCCATTATTTGGGTTTTGTTTAGATCAGAACTTTGCAA  
TGAACCTTTTCGTGCATGGCTGGTCAGTGTCTAAACCATGCTCTGTAGATAAAGTTTCTAAGCCTTGAGGTTGCCTTTCCAGACTTGTGTCAGCTGACGAGGGTAGAGTTCTGCTAC

Primer 1 = GGCTGCTTTTGCACCTGAAAC    Primer 2 = CCAGCCATGCACGAAAAGTTC    Probe = AGACACATCGGTGAGTCGGTGGTTT

### Example of a CR-LOA-wt1 assay (FAM labelled probe)

GGGGCTTGAAGGAGATTTCCCCCAAGCCAC TGCTTGCTTCTGTAGTGAAATGGTCCCTGGGATATAGAGCTGTGGAGTCTCTGA AACTCTGAGCCTCCTGGGAGGAAGAGGGGA  
GACCTGGAATGAGGCCAAGATCAGGTGTGACTTCAAGACTGATGCTTGGTTCTCCGTCTGTACCTCACTTCTTAGGATGTCCCCACCAAGCTTGTGGCCAAGGCCGTGCCTCTGCC  
CATGACGGTGAGAGGCCACTGGTTCCTGAGTCCCCGGACAGAGTACAGTGTGGCAGTGCAGACTGCGGTGAAGCAGAGTGTGGAGAGTACCTGGTGTCTGGCTGGAGTGAGACTG

Primer 1 = TGCTTGCTTCTGTAGTGAAATGGT    Primer 2 = CCCAAGGAGGCTCAGAGTT    Probe = ACCTGGGATATAGAGCTGTGGAGTCTC

**All qPCR assays are run in duplex with a VIC labelled internal control, Dot1l**

### Dot1l internal control (VIC labelled internal control)

AACCAGTGGGCAGTGGATGTGAAGGGCAGGAAGCTCATAGGGTGACTGGCCAACCCAGGGAAGCCGGAGTGCTGCGTCTTCTGTTTCTTCTGTTTCTTTCCCTCTAGTCGTTTTCT  
GTTAGTAGTTGGCATCCTTATGCTTCATCTTACAGTCCGACTTGAGAGCTGCCCCGTAATGGTCGTGCTGGGGCAAGGCTTTATTTCCAGGCGTAGCACACATGGTGGCCAATGGGAC  
TCTGTAGGATCTGCCACACCCATCAGGTGTGCAGGGAGACAGAGCTGAGTCAGGCTCCAGCTCTGGGGAATATGTTGAGTCACCACCTCTGTAGGGTGGTTGTGCATCATAGAAC

Primer 1 = GCCCCAGCACGACCATT    Primer 2 = TAGTTGGCATCCTTATGCTTCATC    Probe = CCAGCTCTCAAGTCG



## qPCR genotyping set up

### qPCR master mix

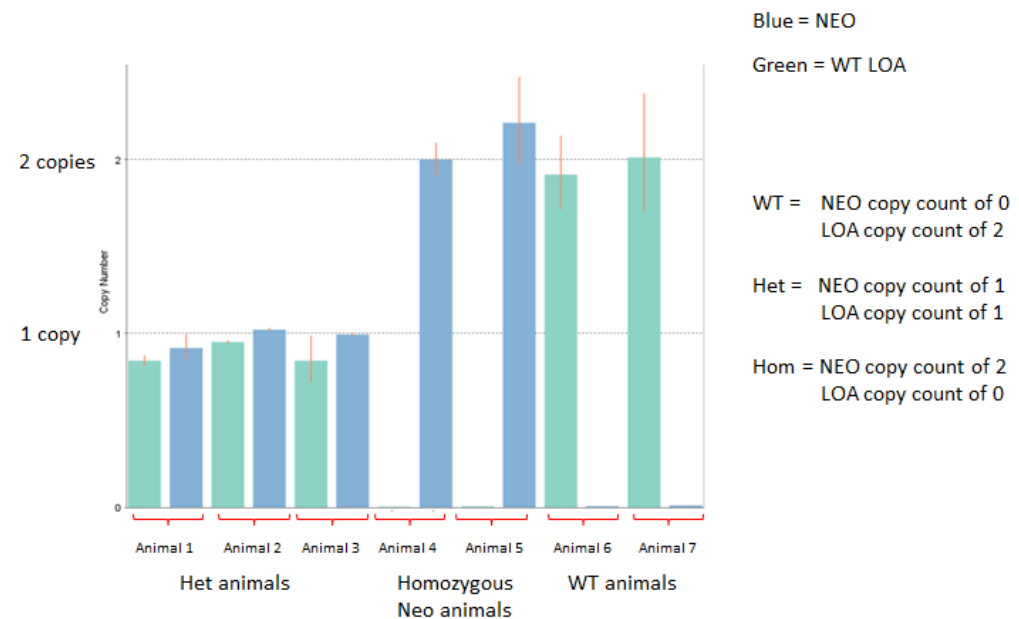
ABI GTX Taqman master mix	5 $\mu$ l
Primers Dot1L_2F (20 $\mu$ M)	0.225 $\mu$ l
Primers Dot1L_R (20 $\mu$ M)	0.225 $\mu$ l
Probe DotL_2M (5 $\mu$ M)	0.2 $\mu$ l
FAM Assay (probe 5 $\mu$ M & primers 15 $\mu$ M each)	0.3 $\mu$ l
Water	1.55 $\mu$ l

### **ALIQUOT 7.5 $\mu$ L**

DNA 2.5 $\mu$ l  
(1/10 dilution of ABI Sample-to-SNP prep)

The results to the right have been run on an ABI 7500 and analysed using the ABI software CopyCaller. Each sample was run with a technical duplicate and for each assay, Neo and the BP-LOA, 7 controls of known copy number and a no template control, all in duplicate, were run so the software can accurately copy count.

### Generic example of a NEO + BP-LOA copy called result





# Pla2g2c

## qPCR based genotyping strategy summary

Assay to use and copy number expected per allele								
	NEO		LACZ		FRT-loxP	Tm1b	BP-LOA	CR-LOA
WT	✓	0					✓ 1	
Tm1a + PL-Tm1a	✓	1					✓ 0	
Tm1b	✓	0	✓	1		✓ 1		✓ 0
PL-Tm1b	✓	1	✓	1				✓ 0
Tm1c					✓	1	✓	0
Tm1d					✓	1	✓	0

Promotor-Tm1b assay is an additional assay that can be run to help identify fully recombined tm1b promotor driven alleles

Expected copy number per animal						
	NEO	LACZ	FRT-loxP	Tm1b	BP-LOA	CR-LOA
WT	0	0	0	0	2	2
Tm1a-HET	1	1	0	0	1	2
Tm1a-HOM	2	2	0	0	0	2
Tm1b-HET	0	1	0	1	1	1
Tm1b-HOM	0	2	0	2	0	0
PL-Tm1b-HET	1	1	0	0	1	1
PL-Tm1b-HOM	2	2	0	0	0	0
Tm1c-HET	0	0	1	0	1	2
Tm1c-HOM	0	0	2	0	0	2
Tm1d-HET	0	0	1	0	1	1
Tm1d-HOM	0	0	2	0	0	0

Assays that are normally run to detect allele

Assays not normally run

Please note copy numbers are based on what is expected for autosomal genes



## Allele conversions

The action of either Flp or Cre recombinase can be used to convert the KO 1<sup>st</sup> alleles described in this document. Both gel based and qPCR genotyping methods may be employed to detect animals carrying the DNA sequences coding for these proteins.

### Cre genotyping

Cre gel based assays are run using primers specific to DNA sequences encoding Cre recombinase as well as a set of primers detecting a house keeping gene Slc40a1. These are run using the same cycling conditions as described in page 4.

#### ActinB-Cre gel based primers

Cre_F1	GCGGTCTGGCAGTAAAACTATC
Cre_R1	GTGAAACAGCATTGCTGCTCACTT
Slc40a1_sh_F (control)	CCTTTGTAACCTCCTCTGTGTC
Slc40a1_sh_R (control)	CTGAAGTCTTTCATGATAACTGCATT

#### Fragment sequences

Slc40a1 147bp

CCTTTGTAACCTCCTCTGTGTC TTTATTTTAGCCTAAACAGCAAAGACTTAAAAGATGGATCTTATCTGGAGAATAGTTCTATTTAGTCCAGCAAACCTTCTAGTAAATATGTTAG  
ATGAAAATGCAGTTATCATGAAAGACTTCAG

Cre1 102bp

GCGGTCTGGCAGTAAAACTATC CAGCAACATTTGGCCAGCTAAACATGCTTCATCGTCGGTCCGGCTGCCACGACCA AGTGACAGCAATGCTGTTTCAC



## Allele conversions

### Cre genotyping

#### Cre qPCR assay (FAM labelled probe)

ATGTCCAATTTACTGACCGTACACCAAATTTGCCTGCATTACCGGTTCGATGCAACGAGTGATGAGGTT **CGCAAGAACCTGATGGACATG** **TTCAGGGATCGCCAGGCGTTT** TCTGA  
GCATACCTGGAAAATGC **TTCGTCCGTTTGCCGGT** CGTGGCGGCATGGTGAAGTTGAATAACCGGAAATGGTTTCCCGCAGAACCTGAAGATGTTTCGCGATTATCTTCTATATC  
TTCAGGCGCGCGGTCTGGCAGTAAAACTATCCAGCAACATTTGGGCCAGCTAAACATGCTTCATCGTTCGGTCCGGGTGCCACGACCAAGTGACAGCAATGCTGTTTCACTGGTT

Primer 1 = **CGCAAGAACCTGATGGACATG**

Primer 2 = **ACCGCAAACGGACAGAA**

Probe = **TTCAGGGATCGCCAGGCGTTT**

qPCR set up for the Cre assay is exactly the same as previously, where the assay is run in duplex with a VIC labelled Dot1l internal control.



## Allele conversions

### Flpe genotyping

Flpe gel based assays are run using primers specific to DNA sequences encoding Flpe recombinase as well as a set of primers detecting a house keeping gene Slc40a1. These are run using the same cycling conditions as described in page 4.

### Flpe gel based primers

ICS_Flpe_F	TCTTTAGCGCAAGGGGTAGGATCG
ICS_Flpe_R	GTCCTGGCCACGGCAGAAGC
Slc40a1_sh_F (control)	CCTTTGTAACCTCCTCTGTGTC
Slc40a1_sh_R (control)	CTGAAGTCTTTCATGATAACTGCATT

### Fragment sequences

Slc40a1 147bp

CCTTTGTAACCTCCTCTGTGTC TTTATTTTAGCCTAAACAGCAAAGACTTAAAAGATGGATCTTATCTGGAGAAtagttctatttagtcCAGCAAACCTTCTAGTAAATATGTTAG  
ATGAA AATGCAGTTATCATGAAAGACTTCAG

ICS-Flpe1 332bp

TCTTTCTTTAGCGCAAGGGGTAGGATCG ATCCACTTGTATATTTGGATGAATTTTTGAGGAATTCTGAACCAGTCCTAAAACGAGTAAATAGGACCGGCAATTCTTCAAGCAACAA  
ACAGGAATACCAATTATTTAAAAGATAACTTAGTCAGATCGTACAACAAGGCTTTGAAGAAAAATGCGCCTTATCCAATCTTTGCTATAAAGAATGGCCAAAATCTCACATTGGAA  
GACATTTGATGACCTCATTCTGTCAATGAAGGGCCTAACGGAGTTGACTAATGTTGTGGAAATTGGAGCGATAAGCGT GCTTCTGCCGTGGCCAGGAC



## Allele conversions

### Flpe genotyping

#### Flpe qPCR assay (FAM labelled probe)

```
GGTAGGATCGATCCACTTGTATATTTGGATGAATTTTTGAGGAATTCTGAACCAGTCCAAAACGAGTAAATAGGACCGGCAATTCCTCAAGCAACAAACAGGAATACCAATTATT  
AAAAGATAACTTAGTCAGATCGTACAACAAGGCTTTGAAGAAAAATGCGCCTTATCCAATCTTTGCTATAAAGAATGGCCAAAATCTCACATTGGAAGACATTTGATGACCTCAT  
TTCTGTCAATGAAGGGCCTAACGGAGTTGACTAATGTTGTGGGAAATTGGAGCGATAAGCGTGCTTCTGCCGTGGCCAGGACAACGTATACTCATCAGATAACAGCAATACCTGAT  
CACTACTTCGCACTAGTTTCTCGGTACTATGCATATGATCCAATATCAAAGGAAATGATAGCATTGAAGGATGAGACTAATCCAATTGAGGAGTGGCAGCATATAGAACAGCTAAA  
GGGTAGTGCTGAAGGAAGCATACGATACCCCGCATGGAATGGGATAATATCACAGGAGGTACTAGACTACCTTTCATCCTACATAAATAGACGCATATAAT
```

Primer 1 = GTGGGAAATTGGAGCGATAAGC

Primer 2 = ACCGAGAACTAGTGCGAAGTAG

Probe = CTTCTGCCGTGGCCAGGACAAC

qPCR set up for the Flpe assay is exactly the same as previously, where the assay is run in duplex with a VIC labelled Dot1l internal control.

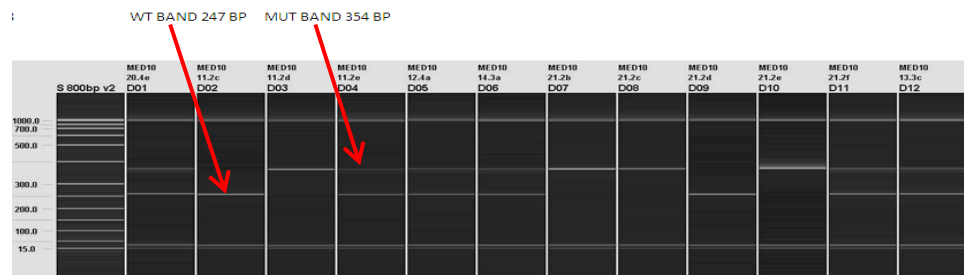




## Lac Z Genotyping Strategy

### Introduction

The gel based assays are normally run on the Qiagen QIAxcel. This is a capillary based system that provides clearer resolution and is quicker than running standard agarose gels. Different size ladders maybe loaded onto runs depending on the fragment sizes being analysed. Typically samples are run with a 50-800bp size ladder.



PCR is performed using KAPA2G Fast Taq polymerase, although alternatives may be used.

### Lac Z gel based primers

LacZ\_2\_small\_F                      ATCACGACGCGCTGTATC  
LacZ\_2\_small\_R                      ACATCGGGCAAATAATATCG

### PCR mix

KAPA Fast Taq PCR master mix	5µl
LacZ_2_small_F	1ul
LacZ_2_small_R	1ul
H <sub>2</sub> O	2µl
DNA	1µl

### Cycling conditions

#### **60TM35FA-(PCR programme specific for KAPA2G Fast mix)**

1. 95°C 1 min.
2. 95°C 10 sec.
3. 60°C 10 sec.
4. 72°C 1 sec.
5. Go to 2 for 34 cycles
6. 72°C 30 sec.
7. 16 °C forever
8. end



## Example of LacZ mutant gel based assay



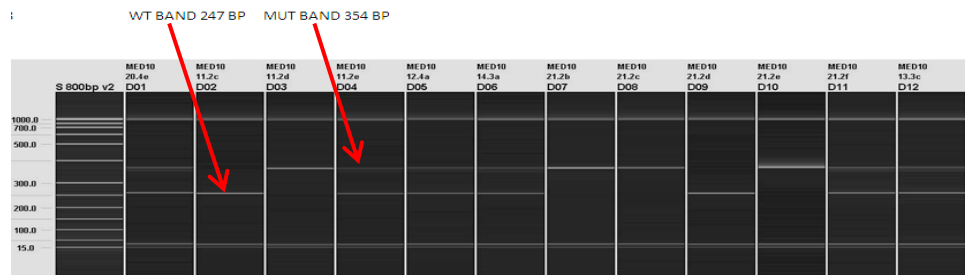
LacZ band = 108bp



## Neo Genotyping Strategy

### Introduction

The gel based assays are normally run on the Qiagen QIAxcel. This is a capillary based system that provides clearer resolution and is quicker than running standard agarose gels. Different size ladders maybe loaded onto runs depending on the fragment sizes being analysed. Typically samples are run with a 50-800bp size ladder.



PCR is performed using KAPA2G Fast Taq polymerase, although alternatives may be used.

### Neo gel based primers

Neo\_long\_F1            TTGAACAAGATGGATTGCACGC  
 Neo\_long\_R1           CCTCGTCCTGCAGTTCATT

### PCR mix

KAPA Fast Taq PCR master mix            5µl  
 Neo\_long\_F1                                    1µl  
 Neo\_long\_R1                                    1µl  
 H<sub>2</sub>O    2µl  
 DNA    1µl

Neo band =186bp

### Cycling conditions

#### **60TM35FA**

1.     95°C    1 min.
2.     95°C    10 sec.
3.     60°C    10 sec.
4.     72°C    1 sec.
5.     Go to 2 for 34 cycles
6.     72°C    30 sec.
7.     16 °C   forever
8.     end