EMMA ID: *02336*
Gene: *Gria2*
Common name: *LEXKO-1605, CHA356C1*
Allele: *Gria2* ^{tm1Lex}

Genotyping Information

Genotyping by end-point PCR based on gel is composed of a genespecific short range PCR using primers on wild type allele and a mutant allele-specific short range PCR. The combined results show the genotype of the mice. For example: mutant positive, wild type positive = Heterozygous.

PCR primer pairs and expected size bands

Assay	Forward Primer	Reverse Primer	Expected Size Band (bp)
Wildtype	LEXKO-1605-16	LEXKO-1605-17	387
Mutant			474
Wildtype (Cre-excised)	LEXKO-1605-F	LEXKO-1605-R	536
Mutant (Cre-excised)			272

Primer sequences

Primer Name	Sequence 5' --> 3'
LEXKO-1605-16	GAGGTGGACATGCTTGGGACAGC
LEXKO-1605-17	CCACATTGATAGCAGTCACCTGC
LEXKO-1605-F	AAGATATGTGATGCAGACACGAGC
LEXKO-1605-R	TCTGCCATTGCTTACTGAGGCTGG

PCR setup (Qiagen, Hot Start Plus)

Component	Volume (µl) 1x	Final conc.
DNA (~ 50-100 ng)	2	
Q-Solution (5x)	2,5	0,5
PCR-Buffer (10x)	2,5	1
DNTP mix (10 mM)	0,5	0,2
$MgCl_2$ (25 mM)	1,5	1,5
Primer 1 (10 pmol/µl)	1	0,4
Primer 2 (10 pmol/µl)	1	0,4
Taq Polymerase (5 U/µl)	0,3	0,06
H_2O*	13,7	
Final volume	25	

*** The amount of H_2O is adjusted with the number of primer.**

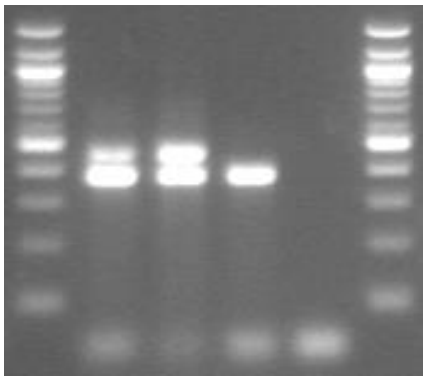
Amplification conditions

PCR Settings	Temperature (°C)	Time	# of cyles
1 Denaturation (Melting)	95°C	5 min	1
2 Amplification (Melting, Annealing, Polym.)	94°C 65-55 (↓1°C/Cycle) 72°C	30 sec 45 sec 45 sec	39
3 Polymerisation	72°C	10 min	1
4 Cooling	4°C	hold	1

use Touch-Down cycling protocol: first 10 cycles anneal at 65°C, decreasing 1°C per cycle, next 30 cycles aneal at 55°C
These PCR conditions have been optimized for our methods and preparation kits. Adaptions may be required.

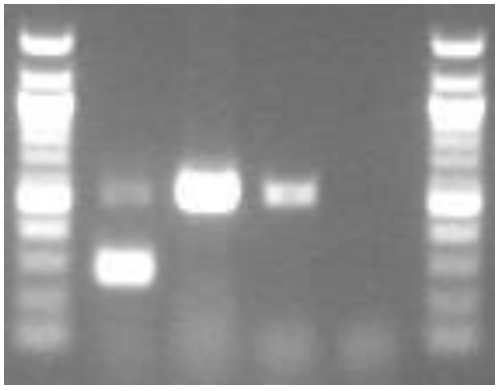
Gel Image

pre Cre



2% AgaroseNEB 100 bp ladder

Cre-excised



2% Agarose, NEB 100 bp ladder