

**EMMA ID:** 02320

**Gene:** *Impa2*

**Common name:** LEXKO-442, PHO252T1

**Allele:** *Impa2*<sup>Gt(OST203987)Lex</sup>

## 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-442-5'	LEXKO-442-3' wt	421
Mutant	LEXKO-442-5'	LTR-rev	135

### Primer sequences

Primer Name	Sequence 5' --> 3'
LEXKO-442-5'	GTCCCTGCGAGGCTAGAATAC
LTR-rev	ATAAACCTCTTGCAGTTGCATC
LEXKO-442-3' wt	GGGCTGCTCTGACGTCTTGG

### 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 <sub>2</sub> (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 <sub>2</sub> O*	13,7	
Final volume	25	

\* The amount of H<sub>2</sub>O is adjusted with the number of primer.

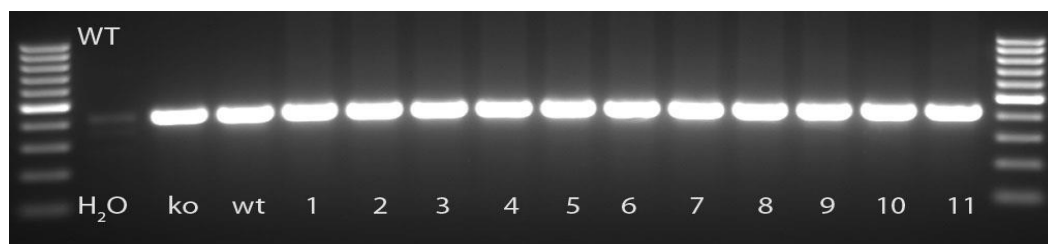
### Amplification conditions

PCR Settings	Temperature (°C)	Time	# of cycles
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

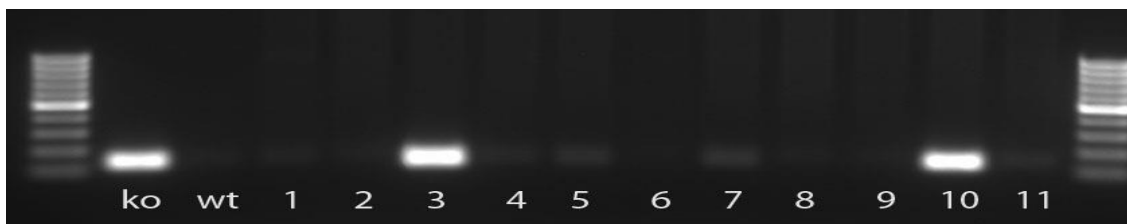
use Touch-Down cycling protocol: first 10 cycles anneal at 65°C, decreasing 1°C per cycle, next 30 cycles anneal at 55°C  
 These PCR conditions have been optimized for our methods and preparation kits. Adaptions may be required.

### Gel Image

#### WT-PCR



#### Mutant-PCR



Separated by gel electrophoresis on a 2% agarose gel.