

**EMMA ID: 05205**

**Gene: *Hprt***

**Common name: *hCMA1***

## 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.

**The *Hprt* gene is x-linked, therfore the males are either hemizygous or wildtype. Females can be homo-, heterozygous, or wildtype.**

**PCR primer pairs and expected size bands**

Assay	Forward Primer	Reverse Primer	Expected Size Band (bp)
Wildtype	-	-	-
Mutant	GX5889	GX2752	2462

**Primer sequences**

Primer Name	Sequence 5' --> 3'
GX5889	TGCTTCAGTCCCATGTTGGCAAGG
GX2752	AAATCTGTGCGGAGCCGAAATCTGG

**PCR setup (Qiagen, Hot Start Plus)**

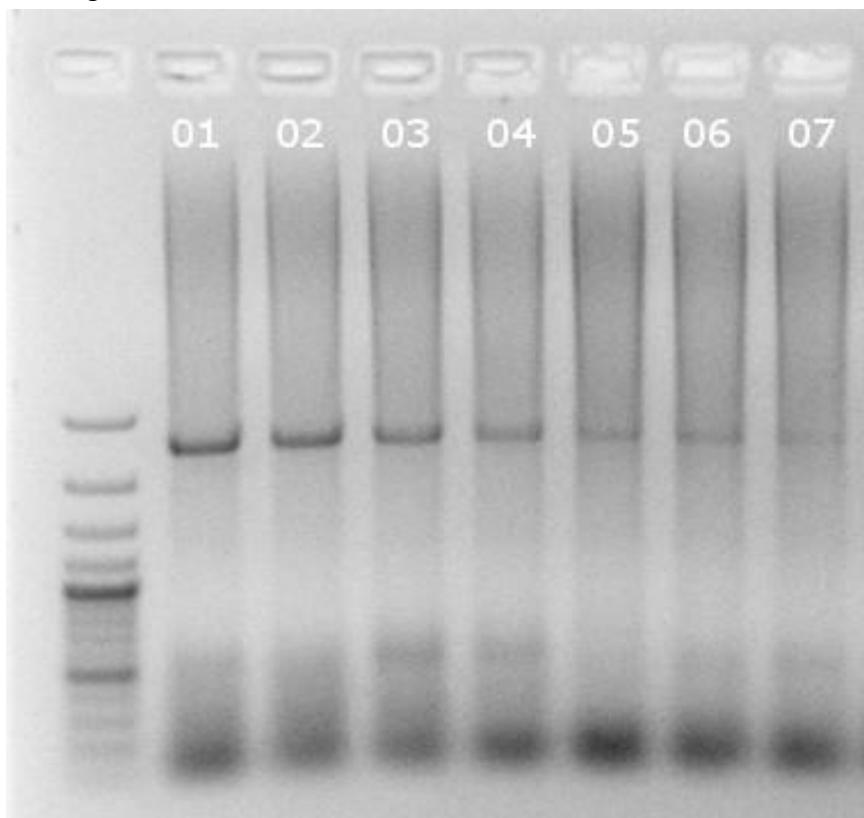
Component	Volume ( $\mu$ l) 1x	Final conc.
DNA (~ 50-100 ng)	2	
Q-Solution (5x)	2,5	0,5
PCR-Buffer (10x)	2,5	1
DNTP mix (10 mM)	0,5	0,2
MgCl <sub>2</sub> (25 mM)	1,5	1,5
Primer 1 (10 pmol/ $\mu$ l)	1	0,4
Primer 2 (10 pmol/ $\mu$ l)	1	0,4
Taq Polymerase (5 U/ $\mu$ l)	0,3	0,06
H <sub>2</sub> O*	13,7	
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°C 72°C	30 sec 45 sec 45 sec	39
3 Polymerisation	72°C	10 min	1
4 Cooling	4°C	hold	1

These PCR conditions have been optimized for our methods and preparation kits. Adoptions may be required.

**Gel Image**


Separated by gel electrophoresis on a 2% agarose gel.